

GENETIC ASSESSMENT OF COLUMBIA RIVER STOCKS

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Executive Summary

This project combines multiple inter-related studies from the Accords Agreement that address the following current and future objectives:

Objective 1) discover and evaluate SNP markers in salmon and steelhead and other anadromous fishes. In the current year of this project we have continued our use of GT-seq protocols for SNP discovery. Our laboratory has designed SNP panels for five study species (Chinook salmon [*Oncorhynchus tshawytscha*] – 299 loci; Steelhead trout [*O. mykiss*] – 390 loci; Sockeye salmon [*O. nerka*] – 382 loci; Coho salmon [*O. kisutch*] – 257 loci; and Pacific lamprey [*Entosphenus tridentata*] – 316 loci). The expanded panel for *O. mykiss* (from 379 to 390 markers) include markers associated with specific traits to enable identification of premature (early-arrival to spawning grounds, primarily summer-run) or mature (late-arrival to spawning grounds, primarily winter-run) steelhead.

In 2019, the total number of samples genotyped with GTseq was 157,319. The largest portion of samples were Chinook salmon (91,516), then Pacific lamprey (26,276), steelhead (23,144), coho (12,310), and sockeye (3,948) as shown in the full section for Objective 1.

Objective 2) expand and create genetic baselines for multiple species including Chinook Salmon (*Oncorhynchus tshawytscha*), steelhead trout (*O. mykiss*), Sockeye Salmon and kokanee (*O. nerka*), and Coho Salmon (*O. kisutch*). Objective two of this project describes efforts to evaluate genetic diversity among populations that will inform managers in the areas of harvest monitoring, and conservation monitoring. Our approach involves the collection, analysis, interpretation and distribution of genotypic data. These data are being compiled as species-specific reference baselines for characterizing Chinook salmon, steelhead trout, and *O. nerka* population structure specific to the Columbia River Basin. The collaborative, inter-agency application of genetic stock identification (GSI) tools continues to provide invaluable monitoring capabilities to understand relative stock proportions in sport, commercial and tribal harvests, as well as monitoring of stock specific run-timing at Bonneville Dam, Lower Granite Dam and other fish weirs in the basin. Moreover, GSI is being used in concert with parentage-based tagging (PBT; *O. tshawytscha*, *O. mykiss*, *O. kisutch*), providing the means to genetically assign individual fish to a hatchery broodstock-of-origin. PBT continues to be valuable for monitoring trends in hatchery production, harvest of hatchery fish, and population attributes of specific hatcheries (e.g., stray rates, survival/mortality, migratory behavior, hatchery/wild interactions). Major accomplishments in 2019 include expansion of our PBT baselines with the addition of several new hatcheries in the Columbia River Basin, both above and below Bonneville Dam. The current PBT baseline for Chinook salmon is shown in Figure 1. The GSI baseline for steelhead trout was substantially expanded 335 SNP markers after genotyping many of the original baseline collections with our revised GT-seq panel.

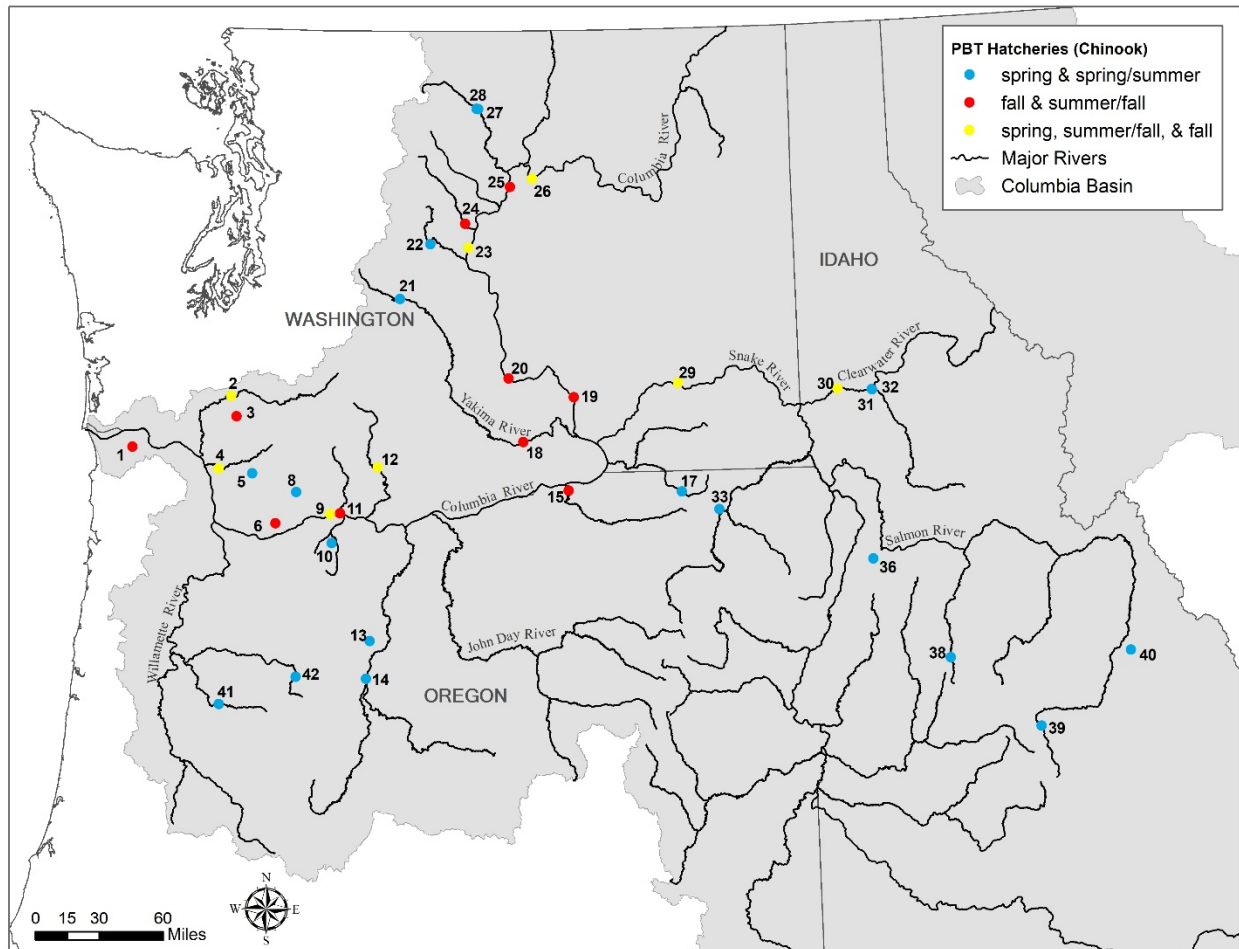


Figure 1. Chinook Salmon, PBT hatcheries. Numbers correspond to map ID and hatchery descriptions.

There are three projects that have begun to characterize reference baselines of millions of SNPs for Chinook Salmon, Coho Salmon, and *O. mykiss*. These new projects utilize whole genome resequencing methods that barcode pools of individuals that represent different collections and can generate allele frequencies across millions of SNP loci. We expect that in the near future there will be no shortage of genetic markers that can be used for GSI and PBT applications anywhere in the species range of our fish of interest. Specific SNPs may be targeted to improve GSI and PBT by incorporating them into GT-seq panels for high-throughput genotyping. In addition, projects will be able to utilize these SNPs for a number of other applications that involve elucidation of genetic mechanisms underlying fitness traits of interest.

Objective 3) implement GSI programs for mainstem Chinook salmon, Sockeye salmon, and steelhead fisheries. In this section, we first described a new method for estimating abundance from catch estimates and at Bonneville Dam using functions that minimize bias from expansion of PBT tag rates. This method allowed us to take PBT tag rates into account, and accurately estimate attributes of natural-origin fish without being influenced by the attributes of unmarked hatchery-origin fish that were not assigned with PBT. We have implemented this approach in all

our harvest estimates for 2018. We have switched to a larger panel of SNP loci (254) that are a subset of the most informative loci for PBT applications which we previously identified among a set of 299 SNPs in Chinook salmon. Results have demonstrated this subset of 254 SNPs improves accuracy of PBT assignments for all lineages of Chinook salmon and effectively avoids crashing issues with SNPPIT, the software we use for PBT assignments.

We used a combination of PBT and GSI analyses to determine stock composition of Chinook salmon harvested in 2018 in test, sport, commercial, and Treaty fisheries in the mainstem Columbia River, and used PBT and GSI to estimate stock composition of sockeye salmon harvested above and below Bonneville Dam in commercial, sport, and Treaty fisheries during the spring, summer, and fall management periods (Figure 2). PBT is a new application for Sockeye salmon and can identify fish that are part of the Yakima River reintroduction using a baseline of translocated adults. This was the first year we analyzed the fall test fishery (pound net) conducted in 2018.

There were 101 coded-wire tags (CWTs) recovered and identified to hatchery stock and broodyear (BY) among the snouts recovered from the lower river fisheries, and 77 of these CWTs also were PBT assigned (76%). Of the 77 fish with both CWT and PBT, there were 65 fish (84%) that appeared concordant with the PBT assignments according to both the hatchery source and the broodyear.

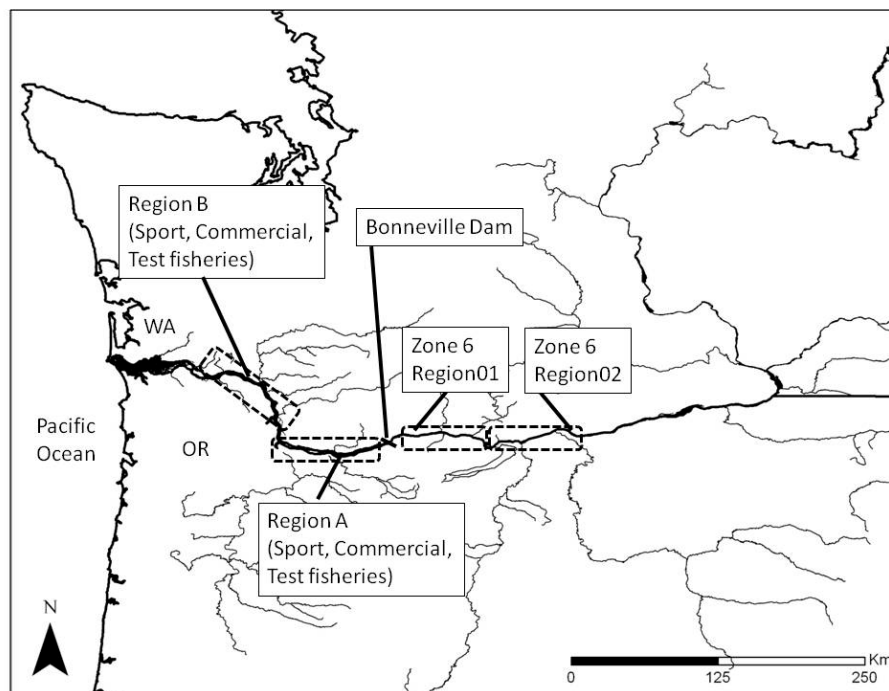


Figure 2. Sources of fishery mixtures in the lower Columbia River mainstem.

We examined one source of information that could potentially be useful to managers particularly on years when the spring Chinook Salmon run is delayed. There is a test fishery that is typically conducted by WDFW on Sundays each week in the early spring (February – May). If the test fishery CPUE were lagged 13 days the peak in CPUE appears to correspond with a peak in Bonneville weekly counts (Figure 3). So if the in-season genetic analysis includes the data

from the test fishery, our report may be able to predict the abundance of particular hatchery and natural-origin stocks that we can expect to pass Bonneville Dam up to 2 weeks later which would be useful information for USvOR managers.

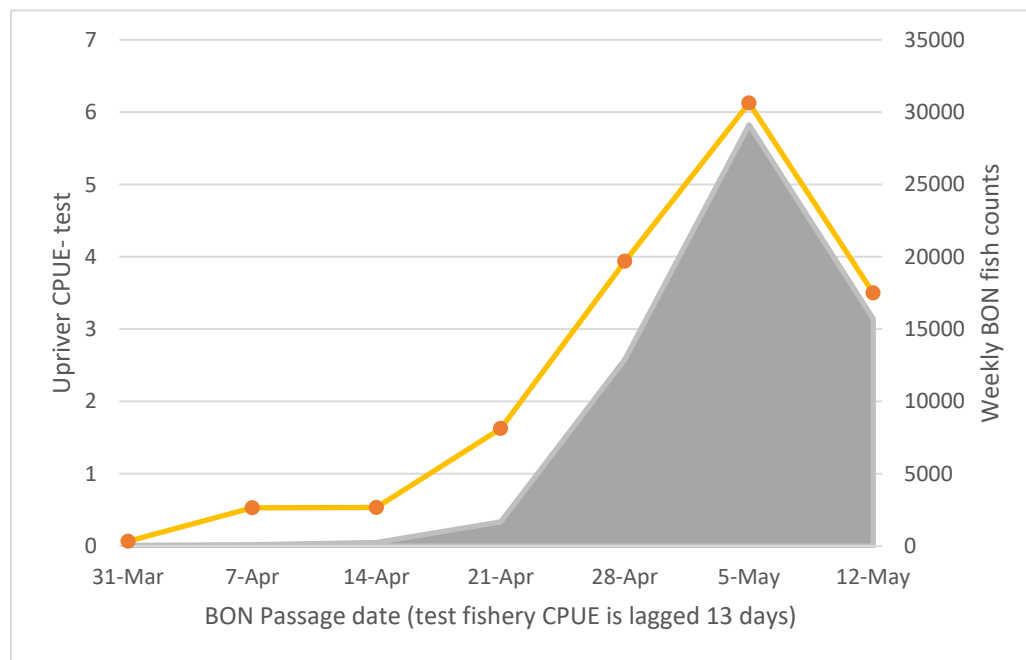


Figure 3. The relationship between the test fishery upriver Chinook Salmon CPUE (line) and weekly fish counts at Bonneville Dam (solid gray) in 2018.

Analysis of Chinook salmon fisheries in the summer management period (June 16 – August 1) addressed the following objectives: 1) estimate stock composition for the mark selective sport fishery in Region B and Region A below Bonneville Dam, and 2) compare stock composition of adipose-clipped versus adipose-intact fish from the Tribal Zone 6 harvest above Bonneville Dam.

The largest difference in the composition of these two fisheries is the presence of lower river stocks (04_WILLAM) in the lower river fishery compared to the zone 6 fishery (**Figure 4**). There were small but detectable abundances of Snake River spring stocks in both the Non-Treaty and Treaty summer fisheries.

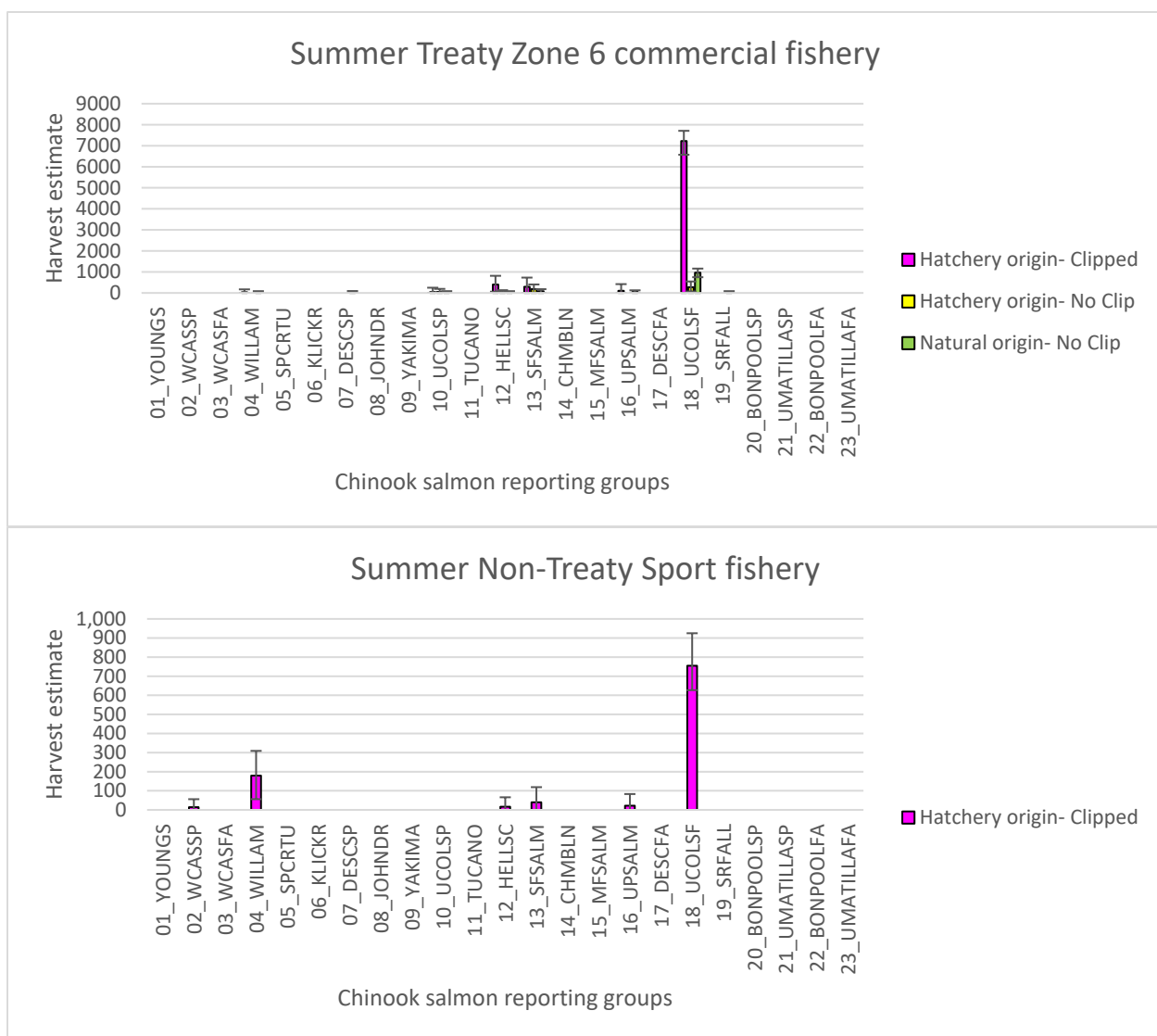


Figure 4. Genetic stock composition of the summer Chinook salmon fisheries analyzed in 2018.

The fall test fishery (pound net) may provide a relatively representative composition of the stocks present in the lower river. Comparison of the test fishery composition and the compositions of the commercial and sport harvests showed a dramatically lower abundance of “tules” in sport fishery compared to the other harvests (**Figure 5**). This difference may largely be due to sport fishers preferentially keeping the VSI bright fish over the tule fish.

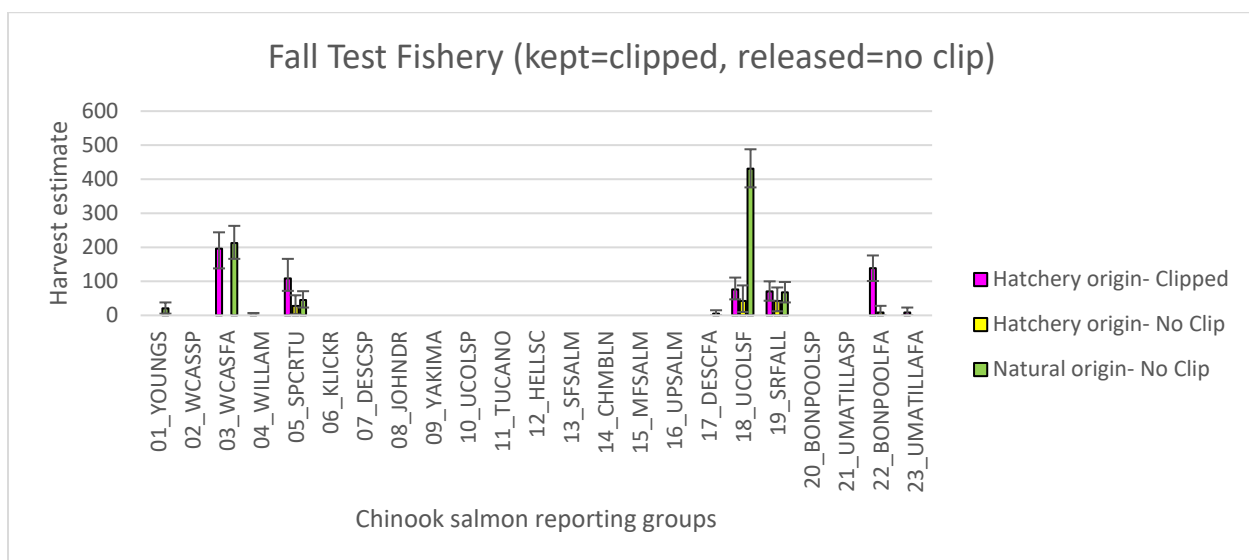
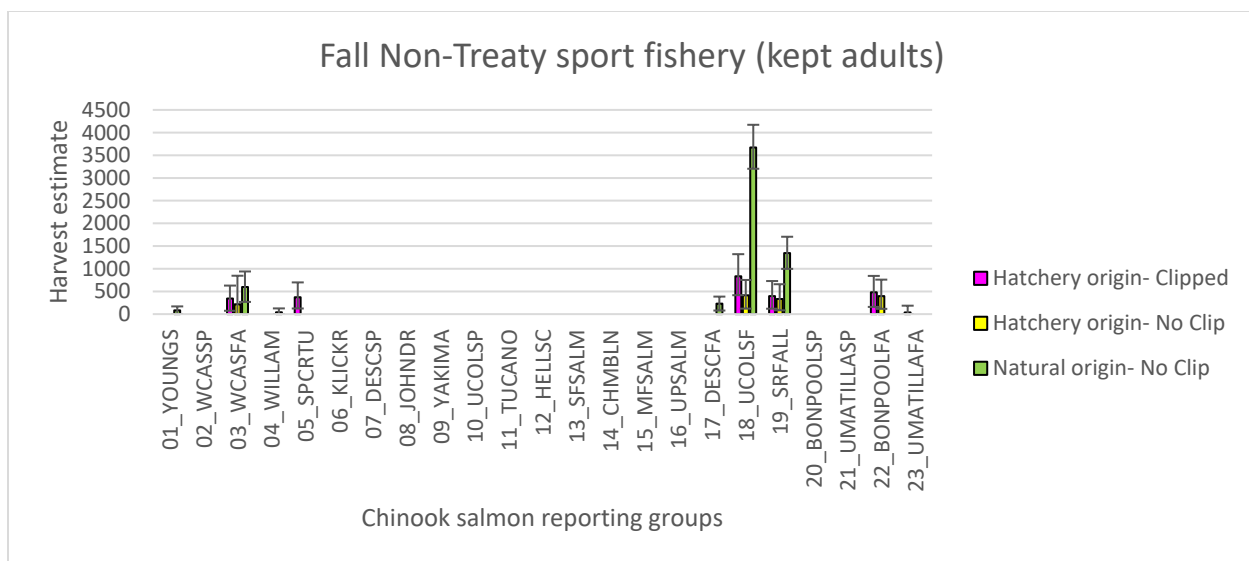


Figure 5. Genetic stock composition of the Non-Treaty fall test and sport Chinook salmon fisheries analyzed in 2018.

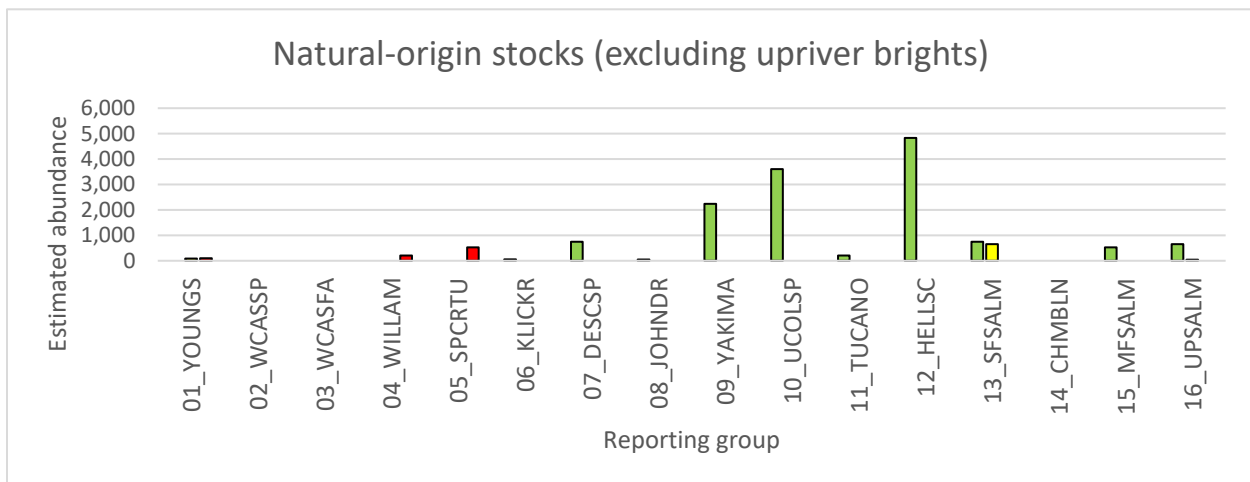
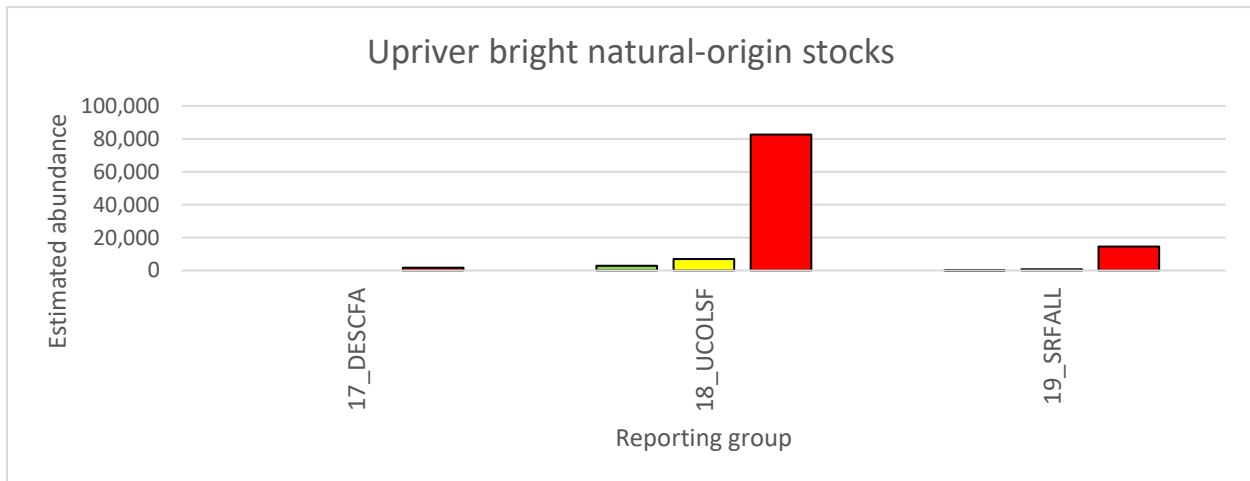
Analysis of stock composition of sockeye salmon fisheries included those from the lower Columbia River sport fishery below Bonneville Dam and the Zone 6 tribal fishery collected in 2018. The overall stock composition of sockeye salmon in these fisheries are shown in Table 1. The proportion of sockeye salmon stocks encountered at Bonneville Dam and in the Zone 6 tribal fishery differed most dramatically for the Wenatchee stock. For the first time, we estimated the number of reintroduced stock from the Yakima River that comprised the fishery (N=149, 2%).

Table 1. Comparison of stock-specific abundance and percent composition among sockeye salmon fisheries. The mean stock abundance estimate is provided for each fishery harvest in 2018.

Mixture source	Mean						Stock proportion					
	Okanogan	Wenatchee	Sna ke	L B C	Yaki ma	oth er	Okanogan	Wenatchee	Sna ke	LB C	Yaki ma	oth er
Sport	111	-	-	-	-	-	100.0 %	-	-	-	-	-
Treaty Zone 6	6,076	1,312	150	5	149	34	78.6%	17.0%	1.9 %	0.1 %	1.9%	0.4 %
Total Harvest	6,187	1,312	150	5	149	34	78.9%	16.7%	1.9 %	0.1 %	1.9%	0.4 %
Bonneville Dam	174,416	17,675	351	80	1,294	-	90.0%	9.1%	0.2 %	0.0 %	0.7%	-

Objective 4) Use PBT and GSI to estimate stock composition of fish passage at Bonneville Dam (steelhead, Sockeye salmon, and Chinook salmon). This section describes our efforts to determine the relative stock composition, abundance and migration run-timing distributions of hatchery and natural origin Chinook salmon, steelhead, and sockeye salmon passing Bonneville Dam. Fish were sampled as they migrated past Bonneville Dam. We sampled adult and jack Chinook and adult steelhead during the spring, summer, and fall management periods, and used a combination of GSI and PBT to estimate run-timing distributions and relative abundance of hatchery and wild Chinook salmon and steelhead stocks in 2018 (post-season analyses). Further, in-season analyses were completed for fish returning throughout 2019 and provided to regional fisheries managers such as the Technical Advisory Committee (TAC).

There were 7 major (i.e., abundance >1000 fish) Chinook salmon stocks represented in the total estimated relative abundance (N=124,854) of natural origin (i.e., adipose non-clipped fish that did not assign via PBT) Chinook salmon passing Bonneville Dam in 2018 (Figure 6). These non-clipped stocks in order of decreasing magnitude were 18_UCOLSF (92,425), 19_SRFALL (15,406), 12_HELLSC (4,833), 10_UCOLSP (3,607), 09_YAKIMA (2,240), 17_DESCFA (1,752), and 13_SFSALM (1,402). These stock abundance estimates were based on the stock proportions that were estimated in using SCOBIDEUX and SPIBETR functions and the estimates of clipped and unclipped adults distributed by TAC.



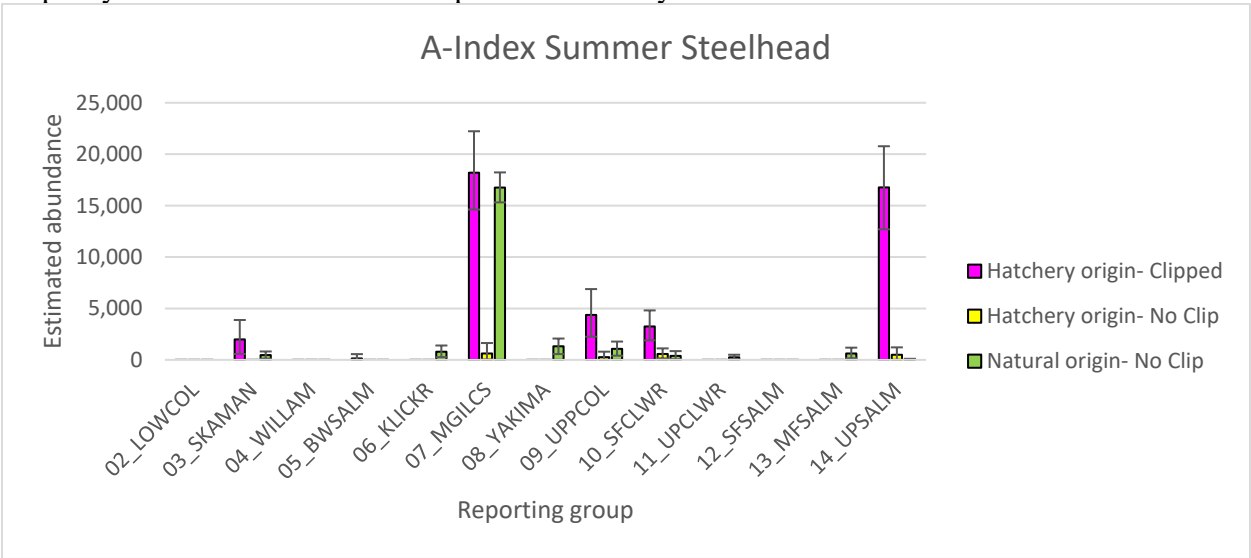
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Figure 6. Estimated abundance of natural origin (excluding adipose-intact hatchery-origin fish) Chinook sampled at Bonneville Dam in 2018 assigned to genetic stock of origin. Upriver bright Chinook salmon reporting groups (top panel), and all other natural-origin Chinook reporting groups (bottom panel) are shown by spring (green), summer (yellow) and fall (red) management period.

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There were five major stocks (abundance >1000) represented in the total estimated relative abundance (N=70,265) of hatchery origin steelhead passing Bonneville Dam in 2018. These stocks in order of decreasing magnitude were 10_SFCLWR (26,807), 07_MGILCS (18,878), 14_UPSALM (17,707), 09_UPPCOL (4,637), and 03_SKAMAN (2,101) (Figure 7). There were three major stocks (abundance >1000) represented in the total estimated relative abundance (N=23,735) of natural origin (excluding adipose unclipped hatchery-origin fish) steelhead passing Bonneville Dam in 2018 (Figure 7). These stocks in order of decreasing magnitude were 07_MGILCS (16,797), 09_UPPCOL (1,424), and 08_YAKIMA (1,321). Two major stocks in 2017 were no longer estimated with abundance >1000: 14_UPSALM (35, down from 2,286), and 06_KLICKR (942, down from 1,013). The 14_UPSALM stock appeared to have decreased dramatically but we feel this was a results of the SCOBIDEUX SPIBETR function that minimized bias from tag rate expansion of the unclipped hatchery-origin fish. This results underscores the importance of the SPIBETR method and using this function consistently on past years of data for better comparisons across years.



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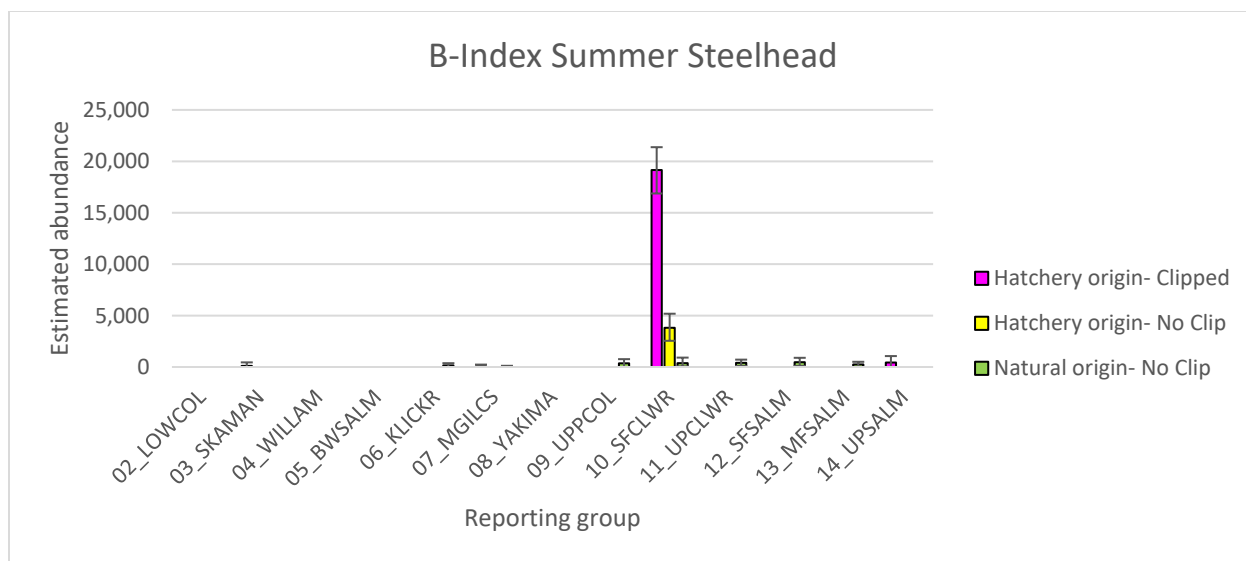


Figure 7. Estimated abundance (\pm 95% CI) of A-Index (<780mm FL, top) and B-Index (\geq 780mm FL, bottom) hatchery origin (adipose clipped and unclipped) and natural-origin steelhead assigned to genetic stock of origin that were sampled at Bonneville Dam in 2018.

Stock abundance for sockeye salmon was estimated over a course of 16 statistical weeks (i.e. weeks 21-36). A total of 1,857 sockeye salmon were sampled at Bonneville Dam in 2018 and were assigned to one of four genetic stocks (i.e., Okanogan, Wenatchee, Snake, and Lake Billy Chinook) using GSI and one reintroduced stock using PBT. The Okanogan stock had the highest relative abundance (174,130), followed by the Wenatchee (17,675). The Snake and Lake Billy Chinook stocks had estimated abundances < 500, but were based on relatively few genetic assignments (<15) (Figure 8). The reintroduced stock from Yakima River was also low sample size but we estimated 1,294 fish in 2018.

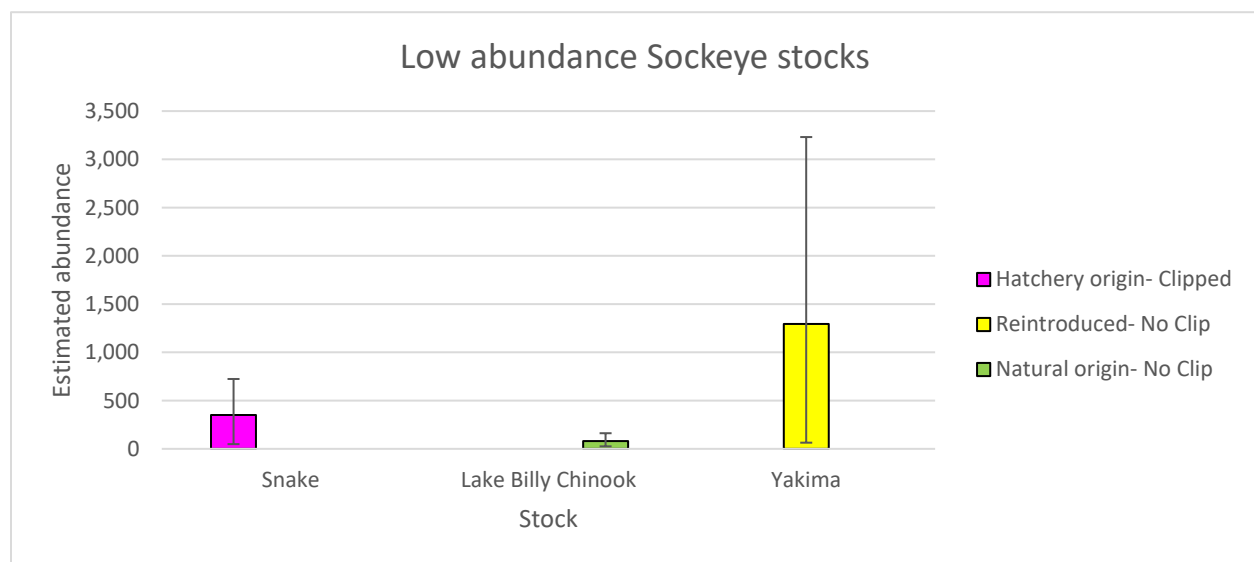
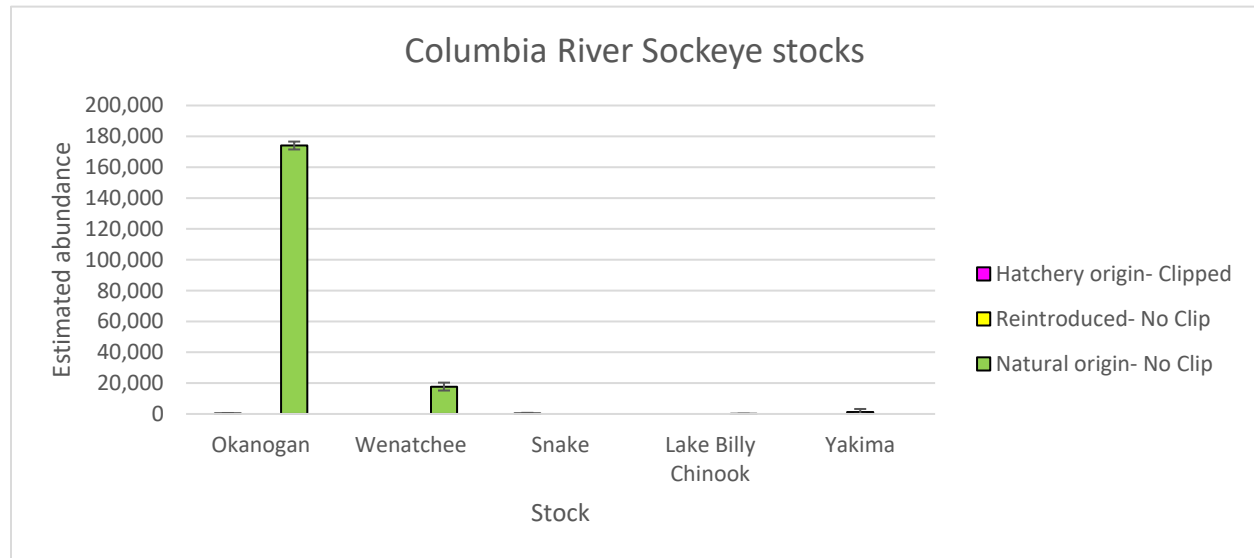


Figure 8. Estimated abundance (\pm 95% CI) of sockeye salmon stocks sampled at Bonneville Dam in 2018.

While the run timing distributions of some hatchery origin and natural origin spring Chinook salmon stocks terminated within the spring management period, several spring Chinook salmon stocks extended well into the summer management period. For steelhead, the patterns generally are consistent with past years. The late arriving natural-origin stocks with median dates on or after August 25th were 10_SFCLWR 11_UPCLWR, 13_MFSALM, and 14_UPSALM. We characterized run-timing by A-Index and B-Index categories for these stocks. For some stocks that had earlier run-timing as A-Index sizes, these stocks were later arriving as B-Index stocks. Run timing distributions for sockeye salmon sampled at Bonneville Dam broadly

overlapped in 2018, and we observed nearly identical run timing distributions for the Okanogan and Wenatchee stocks (median date near 06/24/18).

Finally, we delivered reports to the U.S. v OR Technical Advisory Committee (TAC) on in-season and timely post-season analyses of several runs and fish species passing Bonneville Dam in 2019 (Table 2). The timely reporting of these preliminary analyses increased their usefulness to the fisheries managers of these Columbia River stocks because the results were available while the fishing seasons were being actively shaped or just prior to the TAC needing information to perform their annual analytical tasks.

Table 2. The in-season and post-season report timing and scope of the 2019 fish runs.

Species	Management Period	Data coverage	Analysis begins	Report timing
Chinook	Spring	01/01/2019 – 05/03/2019	5/6/2019	5/10/2019
		01/01/2019 – 05/17/2019	5/20/2019	5/24/2019
		01/01/2019 – 06/01/2019	6/3/2019	6/7/2019
		01/01/2019 – 06/15/2019	6/17/2019	6/21/2019
	Summer	01/01/2019 – 07/05/2019	7/8/2019	7/12/2019
		01/01/2019 – 07/31/2019	8/5/2019	8/9/2019
	Fall	01/01/2019 – 10/25/2019	10/28/2019	11/1/2019
Steelhead	Skamania	04/01/2019 – 06/30/2019	7/8/2019	7/12/2019
	Summer A-/B-Index	07/01/2019 – 07/31/2019	8/5/2019	8/9/2019
		07/01/2019 – 10/25/2019	10/28/2019	11/1/2019
Sockeye	Total	01/01/2019 – 08/02/2019	8/5/2019	8/9/2019

Note: The data were reported as cumulative abundance estimates for each genetic stock during the Chinook Salmon and Summer Steelhead A-/B-Index management periods. The report timing indicates the date these reports were provided to the USvOR TAC chair Stuart Ellis for distribution to TAC members.

Objective 5) Adaptive Genetic Variation associated with environment, landscape, and phenotypic traits

For Objective 5, work has progressed on sequencing Chinook salmon and steelhead throughout the Columbia River Basin to evaluate neutral and adaptive genetic variation related to environmental features. Our recent results indicate that environmental features are strong drivers of adaptive genomic divergence in salmonid species and provide a foundation to investigate how populations might respond to global environmental change (Matala et al. 2014; Hand et al. 2015; Hecht et al. 2015; Micheletti et al. 2018a). Broad geographic patterns of neutral and non-neutral variation demonstrated in these studies can be used to accommodate priorities for regional management and inform long-term conservation of Chinook salmon and steelhead in the Columbia River. Results from these studies indicate that temperature and precipitation are consistently the main environmental factors influencing genetic variation in salmonids, but other variables may also contribute for specific populations. Recent results also emphasize that environmental conditions through the migratory corridor are stronger drivers of selection than those at natal sites for steelhead in the Columbia River Basin (Micheletti et al. 2018a).

Empirical studies have been implemented to advance our understanding of multiple traits related to recovery of salmonids in the Columbia River. Work has focused on genomic regions associated with resident vs. anadromous *O. mykiss* life histories (Narum et al. 2008; Hecht et al. 2013), run-timing related to maturity in steelhead (Hess et al. 2016; Micheletti et al. 2018c) and Chinook salmon (Narum et al. 2018), age at maturity in Chinook salmon (Micheletti and Narum 2018b), disease resistance in *O. mykiss* (Campbell and Narum 2015), and thermally adapted strains of redband trout under heat stress (Narum et al. 2010; Narum et al. 2013; Garvin et al. 2015; Narum et al. 2015, Chen et al. 2018a; Chen et al. 2018b). Candidate markers have been confirmed for adult migration/maturation timing in both steelhead and Chinook salmon and are being screened broadly in large numbers of individuals throughout the Columbia River. Progress has also been made towards investigating the genomic basis for age-at-maturity in steelhead and Chinook salmon, and development of studies to investigate thermal adaptation in anadromous stocks of *O. mykiss* and age/size at maturity in steelhead (A vs. B run). As candidate genes for these traits have begun to be identified, SNP markers from these regions are being incorporated in standard genotyping panels with GTseq in order to validate and monitor genetic variation for these traits in large numbers of individuals.

Objective 6) White Sturgeon Genetics

The research for this project is specific to genetic monitoring of white sturgeon (*Acipenser transmontanus*). The monitoring began in 2008 and has been focused on populations in the impoundments upstream of four middle Columbia River dams: Bonneville, The Dalles, John Day, and McNary. We have also incorporated collections from the lower Columbia River Below Bonneville Dam and from several reaches of the Snake River Basin (Matala et al. 2017). Our monitoring objectives are aimed at providing answers for uncertainties that will inform long-term conservation and management of the species. Some ways this is being achieved is through evaluating the amount of spatial and temporal genetic differentiation among sub-populations or population aggregates to gain a better understanding of the extent of migration (gene flow), relatedness, and effective number of breeders within and between locations. In addition, we are working with the Yakama Nation to genotype supplementation broodstocks utilized for translocation efforts in the middle Columbia but also in several PUDs in the Columbia River above Priest Rapids Dam. Thus far, our analyses suggest there is a limited amount of genetic differentiation between the Middle Columbia impoundments, including hatchery broodstocks. Therefore, hatchery reared fish are not genetically distinct from the wild populations. Isolation by distance indicates a downstream cline in genetic similarity from the upper Snake River down to the Lower Columbia River below Bonneville Dam. This is likely the result of juvenile entrainment at the dams and a lack of upstream migration through fish ladders by adult or larger fish. Recently, we developed a suite of 325 Single Nucleotide Polymorphism (SNP) makers that are being used to evaluate population structure and test the efficacy of the panel in parentage applications. Recent efforts also include progress towards assembly of a draft genome for white sturgeon that will enable further investigation into a sex-linked marker that will allow white sturgeon of any age to be identified by sex. Overall, these efforts are intended to provide an improved understanding of connectivity between adjacent populations (i.e. gene flow), and demographic trends (e.g. age structure) that will be vital in managing for population viability. The contemporary status and temporal trends in genetic diversity are likely to aid managers in understanding the impacts of limited habitat in the confines of the mid-Columbia impoundments that may be exacerbated by further anthropogenic influences.

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Introduction

This project combines multiple inter-related studies from the Fish & Wildlife Program Accords that address the following current and future objectives: 1) discover and evaluate SNP markers in salmon, steelhead, and lamprey; 2) expand and create genetic baselines for multiple species (Chinook, steelhead, sockeye, and coho); 3) implement Genetic Stock Identification (GSI) sampling programs for mainstem Chinook salmon, sockeye salmon, and steelhead fisheries, 4) GSI of fish passing Bonneville Dam (salmon and steelhead), 5) characterize adaptive genetic variation associated with environment, landscape, and phenotypic traits, 6) genetic diversity and structure of white sturgeon. These projects are highly related since SNP markers are needed to complete species-specific baselines, and these baselines are requisite to complete GSI. The results of these four objectives address needs for distinguishing specific stocks, determining genetic diversity, stock specific run timing, and estimating stock composition which can provide information for fisheries management.

Objective 1) SNP Discovery

One of the highest priorities in the full-scale implementation of SNPs for genetic applications of Columbia River fishes is the discovery and development of a sufficient number of markers to characterize population variability. These DNA sequence polymorphisms represent the most abundant variation in the genome of most organisms, and are spread throughout the entire genome at high density (Morin et al. 2004). We currently utilize panels of hundreds of SNP markers in six fish species including: Chinook salmon, Steelhead trout, Sockeye salmon, Coho salmon, Pacific lamprey, and White Sturgeon. Development of these panels of SNP markers has enabled several studies to investigate stock identification of unknown stocks of fish, parentage analysis, population structure, adaptive variation, and underlying genetic variation for specific traits. We also continue to improve our technology for genotyping to make it more cost effective and highly informative to evaluate genetic variation. Our lab developed a new method called GT-seq (Genotyping-in-Thousands by Sequencing) to genotype these SNP markers in large numbers of individuals (Campbell et al. 2015) for much lower costs (< 4 times) than previous technology. Over the past several years, our lab has contributed to the increasing numbers of SNP markers that are available for salmonids and lampreys, and we have reached a point where rigorous stock composition and assessment goals for timely management of fisheries and highly accurate, precise stock assignments can be achieved using panels of SNP markers.

Objective 2) Baseline Expansion

Development and maintenance of genetic baselines have created powerful resources for stock identification of both natural- and hatchery-origin fish. Genetic Stock Identification (GSI) baselines are used to identify the genetic stock of origin of natural-origin fish based on genetic similarity to one of the multiple distinct stocks that represent the Columbia River Basin. Hatchery-origin fish are identified directly to their hatchery of origin based on assignment to hatchery broodstock parents and their ages are inferred using the Parentage Based Tagging (PBT) baselines. PBT continues to be valuable for monitoring trends in hatchery production, harvest of hatchery fish, and population attributes of specific hatcheries (e.g., stray rates, survival/mortality, migratory behavior, hatchery/natural-origin interactions). GSI provides one of

the few ways to estimate the escapement of natural-origin stocks through run reconstruction of these stocks passing Bonneville Dam.

Objectives 3 & 4) Genetic Stock Identification

Genetic Stock Identification (GSI) methods have proven to be effective in determining the proportion of stock origin in several mixed stock applications (Narum et al. 2008b, Hess et al. 2011, Hess and Narum 2011). This study includes two GSI projects that will utilize genetic baselines: 1) GSI to provide information about harvest; and 2) GSI of fish passing Bonneville Dam.

This study includes GSI analysis of Chinook salmon and Sockeye salmon collected from commercial, recreational, and tribal fisheries in the Columbia River. Subsequent years of the study may include other species such as Coho salmon. Implementation of GSI technology could make monitoring individual production units in mixed stock areas possible. Tissues will be sampled annually from fisheries with existing programs in place with Washington Department of Fish and Wildlife (WDFW), Oregon Department of Fish and Wildlife (ODFW), Yakama Nation Fisheries Program (YNFP) and Confederated Tribes of the Warm Springs of Oregon (CTWSRO). We plan to genotype representative samples from fisheries of primary interest. The GSI estimates may help fill information gaps on natural-origin stocks.

The second application of GSI analysis in this study includes sampling unknown origin salmon and steelhead at Bonneville Dam for genetic analysis. Samples will be collected over the majority of the run on a weekly basis, and genetic baselines will be utilized to determine the stock composition of these runs. Few studies have been able to determine the extent of overlap among life history types of salmon and steelhead, but GSI of each life history type will allow us to determine the stock composition of the different runs through Bonneville Dam which can be compared to other methods such as using fish that were PIT-tagged as juveniles. Population genetic methods and statistical assignment models have advanced dramatically in recent years, and estimating stock composition is now possible using either Bayesian or Maximum Likelihood methods (Anderson et al. 2008). Therefore, we plan to estimate stock composition of multiple species passing Bonneville Dam and provide this information on a timely basis to fisheries managers in the form of an annual report.

Finally, we continue to utilize a new genetic technology, parentage based tagging (PBT), in combination with GSI to help augment and refine our stock identification results. PBT is an efficient approach for mass-tagging of fish. The method is carried out by first genotyping a set of potential parents which then provides the opportunity to assign a set of genotyped offspring to their true parent pair. PBT is currently being utilized on a broad scale in the Columbia River Basin to tag all Snake River Chinook salmon and steelhead hatchery broodstocks (Steele et al. 2011) and we now have a baseline that includes most Chinook salmon and steelhead hatcheries located above Bonneville Dam. This application has effectively tagged all hatchery Chinook salmon and steelhead above Bonneville Dam starting with the 2012 brood years. When parent pairs of a hatchery fish are identified with PBT, we can provide accurate information including age of the fish and the source hatchery in which its parents were spawned. We can now use PBT in both Chinook salmon and steelhead GSI applications to identify all hatchery-origin fish, and

then we estimate stock-of-origin of all other hatchery fish that were not assigned with PBT (i.e. the few hatcheries not in the PBT baseline) and all natural-origin fish using GSI. In this way PBT and GSI are complimentary, and using them in combination takes full advantage of the strengths of each method, while resolving or minimizing limitations. Exogenous stock transfers by hatcheries have made hatchery-origin fish challenging to assign with GSI and represents a main limitation that is addressed with PBT. Applications of PBT have been initiated in other species such as Pacific lamprey, and are being used to monitor translocations of lamprey throughout the interior of the Columbia River.

Objective 5) Adaptive Genetic Variation

With increasing genomic information available for non-model organisms, single nucleotide polymorphisms (SNPs) have begun to see increased use as genetic markers for population genetic studies (e.g., Morin et al. 2004). These sequence polymorphisms are densely scattered throughout the genome of most organisms, and are commonly observed in both coding and non-coding regions of functional genes making them ideal markers to study adaptive molecular variation (e.g., Akey et al. 2002). In a large suite of SNPs that are distributed across the genome (e.g., Narum et al. 2018), it is possible to utilize both functionally neutral and adaptive markers within a single study. This combination of information provides a powerful approach to study questions in ecological genetics since both demographic processes (i.e., gene flow and genetic drift) and local adaptation (i.e., selection) may be inferred. Thus, genome scans with large numbers of SNP markers (e.g., RAD sequencing, Baird et al. 2008; Pool-seq, Schlotterer et al. 2014) and gene expression (e.g., RNA-seq) approaches may be effective tools for identifying the genetic architecture underlying specific traits such as thermal tolerance, run-timing/maturation, disease resistance, anadromy, and age-at-maturity. Underlying genomic regions for these traits continue to be investigated and once identified, they can be broadly screened in populations throughout the Columbia River Basin to facilitate management for long term conservation and recovery of salmonids.

Objective 6) White Sturgeon Genetics

The research for this project is specific to genetic monitoring of white sturgeon (*Acipenser transmontanus*). The monitoring began in 2008 and has been focused on populations in the impoundments upstream of four middle Columbia River dams: Bonneville, The Dalles, John Day, and McNary. We have also incorporated collections from the lower Columbia River Below Bonneville Dam and from several reaches of the Snake River Basin (Matala et al. 2017). This research project directly addresses the following uncertainties: 1) What approaches to population recovery and habitat restoration are most effective in regaining meta-population structure and diversity that will increase viability of fish and wildlife in the Columbia River Basin?, 2) How do artificial production and supplementation impact the maintenance or restoration of an ecologically functional metapopulation structure?, and 3) What is the relationship between genetic diversity and ecological and evolutionary performance, and to what extent does the loss of stock diversity reduce the fitness, and hence survival rate and resilience of remaining

populations? Our monitoring objectives are aimed at providing answers for these uncertainties that will inform long-term conservation and management of the species.

Report Structure

This report is divided into six sections, one for each of the objectives of the study. The first section reports on SNP discovery efforts and the second section on genotyping SNP markers in Chinook salmon, steelhead, and *O. nerka* to create genetic baselines. The third section contains stock composition estimates of Chinook salmon and Sockeye salmon sampled in mainstem fisheries in 2018. The fourth section includes analysis of run-timing distributions and estimated abundance of adult Chinook salmon, Sockeye salmon, and steelhead stocks migrating over Bonneville Dam in 2018. In addition, we have implemented in-season and timely post-season analysis of the runs of Chinook salmon, Sockeye salmon and steelhead migrating over Bonneville Dam in 2019. These in-season and post-season 2019 results were provided to fisheries managers that participate on the USvOR Technical Advisory Committee and are summarized in the fourth section of this report. The fifth section reports on progress to identify adaptive variation associated with landscape, environment, and phenotypes. The sixth section reports on updates to genetic analyses of white sturgeon.

Section 1: SNP Discovery

Introduction

Population genetic studies examine variation within the genomes of individuals in order to gain insights into the nature of those populations. For instance, genetic similarities among groups of individuals can indicate relatedness, recent population collapse, or barriers to migration. In the context of salmon conservation, population genetics can answer important questions directly related to fisheries management such as stock exploitation rates, effective population size, and rate of return. Other demographic information such as stock abundance estimates can also be made through analysis of samples taken from fish as they enter the Columbia River through genetic stock identification (GSI). These studies require genotype data from a suitably large and informative set of genetic markers for analysis. Likewise, the number of genotyped individuals must be suitably large to provide accurate results.

Next generation sequencing instruments can provide both a means to identify genetic variation and provide a platform for high-throughput sequencing. In the past, we have used methods such as restriction-site associated DNA sequencing (RAD-seq: <https://www.monitoringmethods.org/Method/Details/4144>) to identify and genotype thousands of single nucleotide polymorphisms (SNPs) within and among study populations. Recently reference genome assemblies have become publicly available for both Chinook salmon and steelhead, and these genome resources has enabled whole genome resequencing for investigating adaptive variation across a large portion of the genome (50-80%) in these species (whole genome resequencing: [Whole Genome Resequencing: Poolseq Individual Barcode v1.0](#)). Whole genome resequencing methods include individually barcoded samples, or pools of samples (Pool-seq, [Whole Genome Resequencing: Poolseq Pooled v1.0](#); Schlotterer et al. 2014) depending on the study design. For both methods, sequence data is aligned to the reference genome assembly, and allele frequencies from millions of SNPs are analyzed to detect statistically significant regions of the genome associated with specific traits or adaptation to environmental factors. Putatively neutral regions of the genome are also useful for standard phylogeny and demographic analyses of populations. In most studies, allele frequencies are available for collections but sequencing depth is typically not high enough to provide individual genotypes. However, candidate SNPs may be developed into standard panels with Genotyping in Thousands by sequencing (GT-seq: <https://www.monitoringresources.org/Document/Method/Details/5446>) or other approaches to genotype many individuals to validate trait association, determine inheritance, and estimate linkage disequilibrium. GT-seq is a high throughput method that uses Illumina sequencers to rapidly genotype thousands of individual samples at hundreds of loci for less than ¼ the cost of previously used TaqMan assays (Campbell et al. 2015). Following the development of GT-seq, our laboratory has designed panels for 5 study species (Chinook salmon [*O. tshawytscha*] – 299 loci; Steelhead trout [*O. mykiss*] – 390 loci; Sockeye salmon [*O. nerka*] – 382 loci; Coho salmon [*O. kisutch*] – 257 loci; and Pacific lamprey [*E. tridentata*] – 316 loci). All of these GT-seq panels have been designed to a maximum of 75bp to allow for inexpensive sequencing runs on Illumina NextSeq 500. Recent additions of 11 SNPs were made to increase GT-seq panels for *O. mykiss* to 390 SNPs. The expanded panel for *O. mykiss* will improve our ability to identify specific traits of premature (summer-run) or mature (winter-run) steelhead (Micheletti et al. 2018).

Methods

For new SNP loci added to panels, the program Primer3 (Rozen and Skaletsky 2000) was used to design primers flanking the target SNP locus for inclusion in existing GT-seq panels. (GT-seq: <https://www.monitoringresources.org/Document/Method/Details/5446>) Parameters used for primer design are as follows (product size range: 50-80 bases, optimal annealing temperature: 60°C, primer size range: 18-24 bases, optimal GC content: 50%). The designed primers were then modified by including the Illumina sequencing primer sites. The primers were ordered from IDT (Integrated DNA technologies) at a concentration of 200µM at the 25nmole synthesis scale. Testing was done by combining primers from previous loci for each species that already worked for GT-seq with the newly designed primers. These new primer pools were then used to create test libraries containing 96 samples using the GT-seq protocol (Campbell et al. 2015). Test libraries were “spiked” into an Illumina HiSeq lane with another sequencing library such that each test library produced about 10 million reads of data for analysis. Since the test library uses only a small percentage of the total reads on the flow cell the new library can be sequenced very cheaply. The sequencing reads were analyzed for the presence of significant numbers of hetero-dimers produced in multiplex PCR using custom perl scripts (<https://github.com/GTseq/GTseq-Pipeline/>). Primers producing large numbers of sequencing artifact reads through primer hetero-dimer interactions were flagged and omitted from the next primer mix. Following this step, the primer mix was used for full scale genotyping using GT-seq libraries containing 3,000-4,000 samples for a NextSeq flow cell.

Results

GT-seq primer pools are being used for all high throughput genotyping projects for 5 target species (Steelhead [*O. mykiss*] – 368 SNP loci including a sex determination marker, Chinook [*O. tshawytscha*]-343 SNP loci including a sex determination marker, Coho [*O. kisutch*]-257 SNP loci, Sockeye [*O. nerka*]-382 SNP loci, and Pacific Lamprey [*E. tridentata*]-295 SNP loci). The remaining primer pools remain unchanged from last year’s report but any additional SNPs from Pool-seq data will be under development in the coming year (Paired-end data assemblies, primer design, and testing). Our GT-seq panels have been used to genotype over 157,319 samples as of Dec. 18th in the 2019 calendar year (Figure 9).

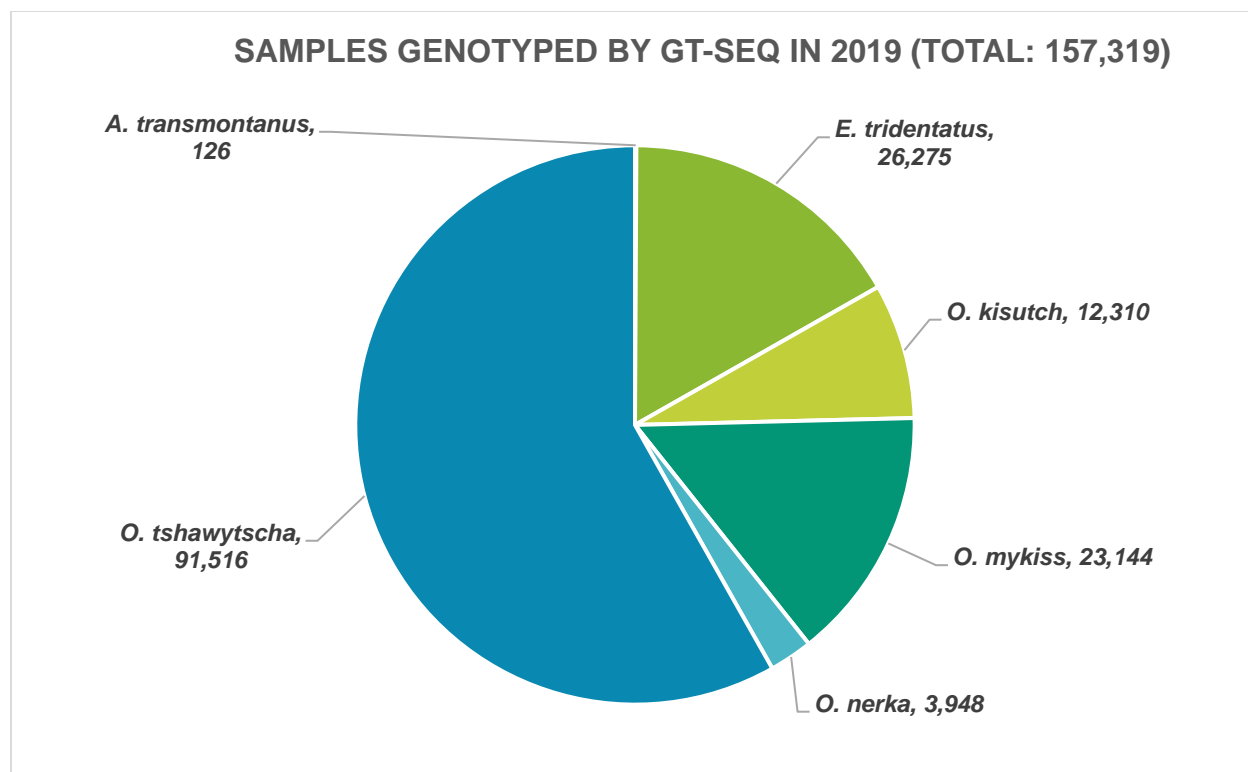


Figure 9. Summary of Columbia River fish samples genotyped using GT-seq in calendar year 2019.

Discussion

The GT-seq genotyping method has allowed for the genotyping of more samples in less time at more loci and at significantly cheaper cost than our previously used method (TaqMan genotyping assays). The total number of samples genotyped using this method has continued to increase from previous years but is expected to stay at a similar level in coming years of this project. The inclusion of more loci afforded by this method has also allowed for improved capabilities such as greater ability to discriminate between reporting groups in GSI and single parent assignments in PBT projects. Similarly, we can now take advantage of genetic markers associated with physical and behavioral traits of our study species by including them in our high-throughput panels. An example of this is our ability to distinguish between early- and late-spawning ground arrival timing in steelhead by including SNP loci found to be highly associated with this trait (Hess et al. 2016, Micheletti et al. 2018).

Expansion of our GT-seq panel for Sockeye salmon is also projected to greatly improve research capabilities in that species. The expanded panel contains 382 loci compared to the previous 93 loci, which should be sufficient for differentiation of Columbia River populations, but also provide statistical power for parentage analysis in this species.

In conclusion, the GT-seq method continues to produce quality genotyping data at a fraction of the cost of previous TaqMan genotyping assays. The technique uses only general laboratory instrumentation (Thermal cyclers, plate centrifuges, quantitative PCR instrument) for library preparation and the Illumina sequencers (either HiSeq or NextSeq500) can be used as high throughput genotyping platforms while maintaining utility for other sequencing studies (whole-genome shotgun, RAD-seq, transcriptome sequencing, synthetic long read, etc.). This is

a key feature of the technique since it allows the multipurpose functionality of the laboratory without investment in specialized equipment. Overall, GT-seq is a valuable tool for conservation genetics studies allowing vastly improved statistical power, higher throughput, and prediction of heritable traits at a lower cost.

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Assay	A	A	Forward	Reverse	A1-Probe	A2-Probe	Allele	GCF_0	GCF_0
	1	2					Correctio	201634	021634
							ns	some	SNP
									Coordinate
OmyY1_2SEX Y	X	Y	GCGCATTTGTATGGTGAAAA	GCCTGGCATATGAGTGT TGA	NA	ATGTGTTCAT ATGCCAG	NA		
Omy_R AD7252 8-44	A	T	TGATGATCCGGACCCTCTCT	CCCGGATTCCCTCCACA GTT	TTGGAACAAA CTGT	TTGGAACATA CTGT	0,0	omy01	7354513
OMS00 078	T	C	GAGGGAAGCAGCCATAAAC AGAATA	GTCTCACTATGGTCCAT ATCTGTGTAGA	TTCACATGCAT AAGAGTG	TCACATGCAT GAGAGTG	0,0	omy01	10499333
Omy_R AD6580 8-68	T	G	TCCTTCACTCTCGATCGGGA	TCAAACCTGGGCCACTAC TGT	ATCGGGATTCT ACTT	ATCGGGAGTCT ACTT	0,0	omy01	12187698
Omy_ga dd45-332	T	C	AGAGAAGACTCACTGCTGTT TGC	AAATCAGTTCCACGCT ATGCT	TTGCTCCAAA ATGG	TTGCTCCGAA ATGG	0,0	omy01	12240053
Omy_10 7031-704	C	T	GGCTTTCGGATACTGAGCAA CAA	TGAACTCACTGTTGGTA TGGACTAGA	TGGACATGAT TGCATAGAC	CTGGACATGA TTACATAGAC	0,0	omy01	18131799
Omy_R AD7396 3-73	T	A	CCCTCTCAGGAAAGTGACCA C	GGATCATGTCAATCTGA TGAGTTGG	TTTCTTTTGGA	TTTCTATTGG A	0,0	omy01	23729539
Omy_m etA-161	T	G	CGCATGCACCAGTTGTAAGA AAG	AGTGCCACCAGCGATAA GAAAA	CAAGTAAGTG GTTATATTCT	CAAGTAAGTG GTTCTATTCT	0,0	omy01	24257340
Omy_R AD5063 2-21	C	T	CCTGCAGGCTGGGTCATTAT	GAGCCAGCTGTACCTTC TCC	TCAGCACCTC CAGCC	TCAGTACCTC CAGCC	0,0	omy01	38986533
Omy_R AD5791 6-29	A	C	GCAGGGCCTTAGAAACAGAC T	TACACGCCTCACTGTTC TGC	CAGGGGCAAAA ACGG	CAGGGGCCAA ACGG	0,0	omy01	44429931
OMS00 070	T	C	CGTTCCTGCGGGACAGT	GTTTCTCTCACGTCCAC AGATCT	CAAAATACGG AAATGCAG	AAATACGGGA ATGCAG	0,0	omy01	55279296
OMS00 003	T	G	GTGCCACTGATGAGGATGAG ATCA	GTAATAAAGCCCTTTTG TGAGGAAAACTAAT	CTTTACTGTCTG ACATTTTA	TACTGTCTGCC ATTTTA	0,0	omy01	59464348

Omy_gd h-271	C	T	AGGTCAGTCTACTTACAGTA TAAAGCAGT	GTCATGTCAACAGAGTA ACATAATAAATCTGC	TCACCCTGAA GTGTAGAC	TCACCCTGAA ATGTAGAC	0,0	omy01	637384 13
Omy_cy p17-153	C	T	GCCCTCCAAGTTCCAAGTGA AAA	CAGGTCATTGATGAAAC GTCAGAAC	ATACCTGAGT GTCATCG	ATACCTGAGT ATCATCG	0,0	omy01	645098 89
OMGH1 PROM1 -SNP1	A	T	TCAAACCTGCATTTGATGGAA ACAAACAT	AGGACAATTCTAAGTGA CCTCAAACCTG	TAGTGTTCACT GACTTCA	TAGTGACAC TGACTTCA	0,0	omy01	704664 74
Omy_G HIP1_2	C	T	TGCATTTGATGGAAACAAAC ATAT-TTATAATGTGT	CAAAAACAAGGACAAT TCTAAGTGACCTC	AAACTGTTGA ACGGTAGTG	AAACTGTTGA ACAGTAGTG	0,0	omy01	704664 83
OMS00 008	A	T	CCCTTTAAGGAGGATTTTAA ATATGTGAGATAGAA	GGATACAGCGTTTTGGA ATGAAACT	CTTCAAATATC CATAATTATAT C	TCAAATATCC ATAATAATAT C	0,0	omy01	773840 35
Omy_10 5385- 406	T	C	GTAACCTACCCTCACCTGAA CTTCA	GTCGCTCTTCTGGGCGT ATCG	CTTGGAACCA TTGCTAC	TTGGAACCGT TGCTAC	0,0	omy01	774461 33
Omy_st at3-273	G	-	CAGACCTCCTCTATCTCCCTA TGAG	ACCTCCTTTAAATTGTG CCCAAGAA	CCAGTTTG	TCAGTTTG	0,0	omy01	79XXX XXX
Omy_G 3PD_2- 371	C	A	GCAGGTAAGGTACACCATAG AGACA	CTCCCCCTGCCTTACCA AAC	AGACATGTGG ATTGGCA	CAGACATGTG TATTGGCA	0,0	omy02	608394 6
Omy_G 3PD_2.2 46	C	T	TCATGTATCAATTAAGGCAT TGTCTTGTCT	GTTAGACACAGTGACCA CCTCTTT	AGTAAAGCCC ATTGTTGAGT	AGTAAAGCCC ATTATTGAGT	0,0	omy02	608407 1
Omy_98 683-165	A	C	GCCATTGCCAGAGAATTTGG TTAA	AACACACGCACCATCTT AAAGC	AGCCAGATAC ATATTTGT	CCAGATACAG ATTTGT	0,0	omy02	855514 0
OMS00 156	A	T	GAGCAGAACACATAGAGGA AAGACT	GTAATCACCTCTTAGC CTGTATGG	TGTGTGTCCTG CTGTAACA	TGTGTCCTGC AGTAACA	0,0	omy02	123971 97
Omy_11 4587- 480	T	G	CAGATTACGTTATTACGTTTG GGAAATTTTAAAGT	GTGAAAGAGTGGGAAA TATAATTATAAGGTCAG A	CCTGTCCAAA ATTGT	CCTGTCCACA ATTGT	0,0	omy02	162045 63
OMS00 138	T	G	TCGGACCACATGAGCAGTTC	GTTCAACAGGTGCCCCAC AC	CTAACAATAA CCAAAGACTG	CTAACAATAA CCACAGACTG	0,0	omy02	223184 48
Omy_N aKATPa 3-50	A	C	GTTGAGCGTGTTATGGGAAA AGAG	TTGCATCGGCTTTCTGA AAACC	CACTCTGTTTC CTTTCTTT	TCTGTTTCCG TTCTTT	0,0	omy02	238931 57
Omy_R AD2504 2-68	G	T	GCTGCTGAAACTGGTTTGCA	TCATGCAGATGAGCTTC CCTG	AATTTCTGCCC AAA	AATTTCTTCC CAA	0,0	omy02	327931 23

Omy_cd 59b-112	C	T	TTTGGATAAGATTGTCTTATA TGACTAAAATGTCATGT	GCCAACGTCCTAGATAT GGTGTAAT	CTAAAAGCCT ATAGCAAACCT	CTAAAAGCCT ATAACAAACCT	0,0,6	omy02	389226 11
Omy_R AD2774 0-55	A	T	TCGGCCTGTACTAGTCTCACT	GCCTAAAATGGCCACTT TCATCA	TAACTTTTAAA AAAA	TAACTTTTAA AAAA	0,0	omy02	427459 63
Omy_u0 9-53,469	T	C	ACAGCCTGAGCGTTTGCA	GGAAACTGGGAGAGAT CAAAGGA	TTGCAGCCCTT ATTGTG	TTGCAGCCCT TGTTGTG	0,0	omy02	488598 83
Omy_R AD2091 7-11	T	C	CGTTGTTCGTCTCCAATCAGG A	ACCAGCTCGATGCCATT GC	AGGTTGCGAG GTC	AGGTCGCGAG GTC	0,0	omy02	523844 97
Omy_m etB-138	T	A	TCTGTCCCTGACGCTATAAA AACG	GAAGTATTTTCAGCTTAA TTTCACTGTTGAGTT	TTCGCCAAAG AGAAAT	TTCGCCAAAG TGAAAT	0,0	omy02	537920 50
Omy_R AD9004 -13	G	A	TATACCACGCCTTCCCTGGA	CAGAGAGAAATCCCCCA CCC	TCATCTGAAG GGGG	TCATCTAAAG GGGG	0,0	omy02	540448 36
Omy_lpl -220	C	G	TGACAATCACTGAGCAACTG AACTC	GTCCAGTCTTGCTTCAA CTCATTCT	AGTGACAGTC A	AGTCACAGTC A	0,0	omy02	600228 13
Omy_R AD1610 4-20	A	G	ATTCCAAAACCTGCAGGGGT	TCAGGATTTGGTAAGGT GGCC	AGGGCAAAG[AT]CAAAGG	AGGGCAAGG[AT]CAAAGG	0,0	omy02	671193 13
Omy_R AD3781 6-68	A	T	CTCATTCCTGGCCGTCTG	CCACTCACACTGGCTTA TGC	GCGGCGTAAA AATG	GCGGCGTTAA AATG	0,0	omy02	735000 31
Omy_R AD4631 4-35	A	G	ACTGCATCTTTTCCCCTGCA	TGAAGATACCCAGAGAC ACCA	TAGCAATGGT	TAGCGATGGT	0,0	omy02	784801 58
Omy_R AD7320 4-63	G	C	CCTGGGCAATGACCTCCAC	AGCTCCCTTCTCTCTCCC TC	GTGCCCCTCT CCACCG	GTGCCCCCTC TCCACCG	0,0	omy03	246702 64
OMS00 096	T	G	CATGAGAATGGATCAGTCTC CACAA	GATGAAATCTGAATGTG TTGACACTACAG	AAAGAGGAAG AGTCTCG	AAAGAGGAA GCGTCTCG	0,0	omy03	291093 94
Omy_11 2301- 202	T	G	GTAAACCCTGCCCACATAAT TAGGT	CTGAGACACTGCTCCAA GGT	AATGCGAAGA CAAACCT	AATGCGAAGC CAAACCT	0,0	omy03	375905 54
Omy_R AD3379 8-24	T	C	CAGGAGGGTCAAGTGGAGTC	TTGGGCCCTCTCTTTTTG GG	GAGTCTATCA AGAT	GAGTCTACCA AGAT	1,4,0	omy03	383366 51

Omy_R AD1349 9-13	T	C	GTTACCTGACGACCAAGGT	GCTGGGGGAGCTTTACA TGA	CGCCCTGTCC GCCA	CGCCCTGCCC GCCA	0,0	omy03	384089 41
Omy_u0 9-54- 311	C	T	GTGGCTCCCCAGGAACAAG	AAGTTTCATGTCACATT CCAGTTACCT	TGGTAATTATT CAACAGATCA GT	TGGTAATTAT TCAACAAATC AGT	0,0	omy03	429205 02
Omy_R AD5883 5-15	G	T	GTCTGCTAAGGTCCTGCAGG	GCCGACCATGAGAGACC TG	ATAGCTGCTG GGACCCA	ATAGCTTCTG GGACCCA	0,0.2	omy03	537353 38
Omy_u0 7-79- 166	G	T	CCCGCTATATTATTTGATCAC CCTTGA	ATTTAAATCCATTTCTA AAAATAAGCAAACCTA ACCA	ACTTGGGAAT ACCCAGCC	CTTGGGAATA ACCCAGCC	0,0	omy03	575236 76
Omy_11 7815-81	C	T	CTGCTTTATGCACACCACATT GT	GCTCTTTCTGGAGAACA AGGTACTG	CTATACGGAG ACCAGC	CTATACGGAA ACCAGC	0,0	omy03	678942 57
Omy_11 8654-91	A	G	CAGCGTAGACCGTTTCCTCA TTAT	GCGCCGATGAGCAGCTT	TCAGCTTGTCT TGCCGC	CAGCTTGTCC TGCCGC	0,0	omy03	754580 69
Omy_ar omat- 280	T	C	CTCCATTGATTTCATGCCGAA CATT	GGAGAGGTCAAACATA GCCTGGTA	TCTTGCAAAC TCC	TCTTGCGAAC TCC	1,0	omy04	339146 9
Omy_10 5075- 162	T	G	GGAGAAGGACAAGGACATT GGTAAT	AAAGCAGACCACACCAT ACTTCTC	CTTTCTCTCCT ACTTTCC	CTTTCTCTCCT CCTTTCC	0,0	omy04	763533 4
Omy_m yoD-178	A	C	GGTCAAATATTTTCATTTACG ATTACACTTAGGC	TGGCAAAGCTGTCATTC CTTCTAAT	TTTTATGAGAT ATAATTTCC	TTTTATGAGA TATCATTTCC	0,0	omy04	105207 93
Omy_12 8923- 433	T	C	CTATGTCCTTGGCAGAAGTC TACA	ACGTTTCTTTGGGCTGA GACTTATT	CTTCATTTTCA TTCCTGTTTT	CATTTTCATT CGCTGTTTT	0,0	omy04	164362 34
Omy_13 0524- 160	C	G	CGAAGGTAGCGATTGGTCGT T	TGTCTGTTCTGCTGTGTG CTT	ATGGCTTGAT CCTCA	ATGGCTTCAT CCTCA	0,0	omy04	288890 24
Omy_R AD7778 9-54	T	C	AGACAAAACCTGCAGGGGA C	AGCACGTTAAAACCAAA CTGTCA	TAAATTATATT TGACAG	TAAATTACAT TTGACAG	0,0	omy04	309761 38
OMS00 079	T	C	GTAACATTATGAATCTATCA GTTTCCCTAGCT	ACCTGCAACGTTAGAGC TGTTTATT	CTACTTTTCAC AGTAACACAG	CTACTTTTCA CAGTGACACA G	0,0	omy04	352504 73
Omy_11 7286- 374	A	T	TGATGTGTTGTTCCCTCATGGC TTA	CTGTGCATTTATTCTTGT GATGCTAGG	CTTTCCTCATC ATACTCTATG G	TCCTCATCAT ACACTATGG	0,0	omy04	523101 65

Oms00 087	A	G	GCAAATTTTACCCTTAACGT GGTTT	GATTTGATGTGTGTGTA TTACCTCCTCTA	GTTA[CA]AAC TGACAAAGTG TG	GTTA[CA]AGC TGACAAAGTG T	0,0	omy04	525119 64
Oms00 111	T	C	CATGCGGACCTGCATAGCT	GCTTAGCCATTGACAGA GCATATCA	CAACCAGACT ACCATTC	AACCAGACTG CCATTC	0,0	omy04	630315 81
Omy_R AD9248 5-64	T	A	CCAGTCAGTCTTGCCTCAGG	GGTCACCACAGGATTGG AGG	GTGTAGATAT ACAT	GTGTAGAAAT ACAT	0,0	omy04	683184 64
Oms00 116	T	A	GCCTTTCTCCCATATCACATT CGA	AAACGCATCTTACACTG TGTTGTG	CTTTTACATTT TCAATATTCTG	TTTACATTTTC AATTTTCTG	0,0	omy05	113417 54
Omy_R AD1307 3-16	G	A	GTGAGGGATCACACCTGCAG	GCACCCATTTCGTAATGT CCC	AAAGGGGACA TTACG	AAAAGGGAC ATTACG	0,0	omy05	139545 98
Omy_11 0362- 585	G	A	GCAGCCAAGATGAACGAAA ACTTC	CCGGCCTGGGTCTCAAT G	CACCGCCCTG CCCGT	CACCGCCTTG CCCGT	0,0	omy05	143072 79
OmyR2 4370	G	A	TCATTACCTACGCAGTGGAG	ATCTCTGGGCCTGAACA AT	ATTTAGCAGG AGGCCTCTCC A	ATTTAGCAGA AGGCCTCTCC A	0,0	omy05	285793 73
OmyR4 0252	T	A	ACTCTGAATTCCTCAGGCTT	TCGAACCAGCTGTCTTT CT	AATGCTATATT GAACCTTAA	AATGCTAAAT TGAACCTTAA	0,0	omy05	316752 78
OmyR1 9198	T	A	GTAACCATGACATCCACCAA TC	CCTATGCACAAAGCCTT CAC	ATCCCTGTTAT CTAATCATT	ATCCCTGTAA TCTAATCATT	0,0	omy05	349734 85
Omy_R AD2389 4-58	A	T	TGCAGAAAGGCTGTGTGGAT	TCTTAACACAGTCCTCA TGGAACA	GTGGATTAGG GG	GTGGATTTGG GG	0,0	omy05	409271 21
OmyR3 3562	G	A	CAGGAGTAATGCATCCCAAT G	CAAGTTGATAACGAGAC ATAAGGG	TAAGACTTGG CATAAGCATG	TAAGACTTAG CATAAGCATG	0,0	omy05	473375 40
Omy_bc AKala- 380rd	G	A	TTGCTCTCTTCTGGTTGCCTT A	CTTCAGGAGAAAGCGCT ACTGT	CATACCCATC CTATGTCAG	CATACTCATC CTATGTCAG	0,0	omy05	534692 95
OmyR1 4589	A	G	GGTGACTGAGCTGGATGT	TTTAGAGAATTTGGCAG TACGTC	GTGGTTACAA AGGGTCTGCA	GTGGTTACAA GGGGTCTGCA	0,0	omy05	561627 85
Omy_u0 9-61.043	A	T	TAGTCACATCCATAGTAATA CTTCC	TGTTTCAGAAGCAGAAAA CCAATCTCT	CACTTGTCCT TTTTCA	CTTGGTCCAT TTTTCA	0,0.4	omy05	588408 10
Omy_R AD3039 2-17	T	C	CCACTACTCACAGACCTGCA	GCTCAAGGACCAACAA AAAGCT	CTGAGACTGT GTGT	CTGAGACCGT GTGT	0,0	omy05	617726 69

Omy_S ECC22b -88	T	C	GGATCCCTCCTTTTAACACA AGACT	CTACAGGATGACTACCT AATTGCTAATAAAACA	CTGTCTGTCCA TATATC	CTGTCTGTCC GTATATC	0,0	omy05	618289 03
OmyR4 0319	C	T	AAAGATTGCTGCGATGTCTA AT	CGCAGAGAACAGAGGA TGA	TGCAAGTGC[T A]GCCCTTTTA T	TGCAAGTGC[TA]GCCTTTTT AT	0,0	omy05	631589 19
OMS00 169	A	G	AGCACTTGACTCAAACAC ATAAATCA	CTGAGACAGGAAGAAC AATGTTAACAAAA	CAAAAAGCAT TGATATCAAT	AAAAGCATTG ACATCAAT	0,0	omy05	675681 74
Omy_R AD8513 1-35	T	C	TTCAATAACTACAGGCAGAT GGT	AGTTCCCAAATGCACTG TACA	GATGGTATGG TGAG	GATGGTACGG TGAG	0,0	omy05	755346 97
Omy_10 9525- 403	A	G	CCTCATTCTCATTGGTGAGTT GTCT	TGTAAGATCTGACCACA TGAGTATAACCA	CCTACACCTCT TTTTTCCACA	CCTACACCTC TTTTCTCCAC A	0,0	omy05	842248 92
Omy_R AD4357 3-37	A	G	TGCAGGGAACGATGAAACCA	ACAGACACAGCATTGGC CAA	GAAAGAGAGA GTTT	GAAAGAGGG AGTTT	0,0	omy06	174595 4
Omy_10 5714- 265	C	T	CCACTCAGTGCAAGCATGGA	GCTTTCAATCCTTGGCT CCAATATC	CTGTTGTTTGA GGTTCAG	TGTTGTTTGA GATTCAG	0,0	omy06	759484 7
Omy_10 7285-69	C	G	GCCCTTGTGACAATGCACTG TTATA	AGGTCTAGACAGTGTGC CATTTG	ATACGTTACTT TTGACCTTGT	ACGTTACTTT TCACCTTGT	0,0	omy06	980915 6
OMS00 013	A	G	GCCTTTGTTCTCCTTGGTGGT TA	AGAAAAGTGTGGACTG AGGTTGAG	CTTCTTTTCCC TTGCTACTC	CTTTTCCCTC GCTACTC	0,0	omy06	103252 71
Omy_R AD3541 7-9	G	A	GCACTTGACCACATAGCTGG	ACTCCACACTCCACAAA GCA	TGCAGGACGT GCTTTGT	TGCAGGACAT GCTTTGT	0,0	omy06	158675 14
Omy_b9 -164	T	-	GCACAGAACACAGCCAATAT TAACA	GCCTTGACTCTCCCTTC ATGAC	CCTACAACCTT GATCTAACGT G	CCTACAACCTT GATCTACGTG	0,0	omy06	175997 31
Omy_R AD4795 5-51	G	T	AGTGTGCTAGAATGGGCCTG	ACCATGGGCAGTTCATT TCA	TTGGAATAGA ATCTATA	TTGGAATATA ATCTATA	0,0	omy06	231917 48
Omy_va mp5- 303	A	-	CTGCTTCCCAATTCAGTATCG TCTT	AGGCTGAAGCATTTCTG AGTATGAA	TGGCCGTAGT AGTTGGTCA	TGGCCGTAGT TGGTCA	0,0	omy06	336251 38
Omy_st ar-206	A	G	CGTGTGCCAGCCCTTCT	GACCACTGAGATCATTG CTGTGA	TCTTTGGCACT ATATCT	TTTGGCACCA TATCT	0,0	omy06	366248 63

Omy_R AD6863 4-40	A	C	TGCAGGACTCCTTTGAAACG T	TAACGCCAGCTGCATGA TGA	CCTCTAAACT GAAT	CCTCTAACCT GAAT	0,0	omy06	402608 00
Omy_R AD3695 2-53	C	A	TGTACGTCATTGGGGCTGAG	CCTACCAGACCACACGA TGA	AGGACATCTT CATC	AGGACATATT CATC	0,0	omy06	402609 70
Omy_IL 1b-163	T	G	GGAACAACAGGATTAAGCCT ACTCT	CCTAAAGGCCTAGGAAA CTAAACTTCA	CTGAGGTCAT AAAAATA	CTGAGGTCAT ACAAATA	0,0	omy06	423131 29
Omy_R AD7016 -31	C	A	GCAGGAATATTCACTGTTGC CA	TCTAAAATGTCGTTGGC GGC	ATAATTTTATT TAA	ATAATTTAAT TTAA	0,0	omy06	514907 60
Omy_R AD6013 5-12	C	G	AGCATACACACCTGCAGGAA	TGGTAGGAGGAGATGCT CTGT	GAACATACCG GAAC	GAACATAGCG GAAC	0,0	omy06	561100 72
Omy_R AD3514 9-9	G	A	GAGTCAATAGAGCCCCCTGC	TGGTTAGCAGGAGCAAT CTCA	GCGCGCTTAT GTC	GCGCACTTAT GTC	0,0	omy06	601825 01
Omy_R AD3915 6-33	T	C	GGGTGTGACATGTGTGCAGA	ACTGCTTGTCCCCACCA AG	ACCGTAATGG AGAG	ACCGTAACGG AGAG	0,0	omy06	468XX XXX
Omy_R AD366- 7	C	A	ACCAAATTAGAGCCTGCAGG A	GGAGAGGCCTTTCCGTG ATC	CAGGACTTGC TTTT	CAGGAATTGC TTTT	0,1.4	omy07	623006 6
Omy_m apK3- 103	A	T	GAAGTCATTACTGGTCAGTG GTCAA	GCACAAAACATGAGGA AAGTTGAGA	AATTATTAAG CCTATTTTTTT	ATTATTAAGC CTAATTTTTTT	0,0	omy07	109757 16
Omy_O myclmk 438-96	A	C	CCCGACTCTACTTCACTACTT TCCT	GGCCTAGGACAATAGG ACTGAAC	TACGCAAATT AGGTTTAAA	CGCAAATTAG GGTTAAA	0,0	omy07	109760 28
OMS00 176	T	G	GTTGGAAGTTCCGGTGGTAG AG	CTGGGTCCTGAAGGAGC TT	TTCCAGCACT GCTGTC	CCAGCCCTGC TGTC	0,0	omy07	110291 67
Omy_13 1460- 646	C	T	GTGAAAAGGAATGGAGGAG TACAGT	TGCTAGGACAGGAAGAT CATTTGTG	AATAAAGCAG AATTTGTTACT G	AAAGCAGAAT TTATTACTG	0,0	omy07	366000 38
Omy_R AD3061 9-61	T	A	CTGCAGGTCAATGGGTGCTA	ACACTGATCACATTTTT GTCACACT	CACTGTAAAA	CACTGTAAAA	0,0	omy07	427104 19

Omy_R AD2567-8	A	T	CTGTCTGGATAGCCTTGCCC	TCTATCTCTGGGGAAAA TAGCCC	GCAATGGGCT ATTT	GCATTGGGCT ATTT	0,0	omy07	437932 41
Omy_p ad-196	C	T	CAAACAACCACAGTAGTCCT CCAAT	GCTTTTCACCCTTTTGTA AATTAAGCCAAA	AAGACAAAGG TGTAATACC	AAGACAAAG GTATAATACC	0,0	omy07	444706 19
Omy_R AD1243 9-64	G	A	GGAACTTTTCACATCATGTT GACTG	GCACAGAGAACTCCAG GCAA	CTTCTCCGATG TCA	CTTCTCCAAT GTCA	0,0	omy07	445378 93
OMS00 064	T	G	GTGGATATGTAGTTCGATGG AACAGT	TTTACAACAATCTTCTTT TAATAAAAAATATAGCCA CTTAT	CAGGCAACAT TTTATATAACT A	CAGGCAACAT TTTATCTAAC TA	0,0	omy07	452277 50
Omy_R AD6259 6-38	A	T	GCAGGACACTGGTTCCCAAA	CCTGAGATTTGAGATCA CTGGCT	TTAAAAAATA TATATTA	TTAAAATATA TATATTA	0,5,0	omy07	499777 94
OMS00 154	A	T	GATGTTGGCTGGAGGTGTAG T	TGGGAACACTTTGCCTA CCC	ACAGGGCTTC TGATTGA	AGGGCTTCAG ATTGA	0,0	omy07	562345 73
Omy_sy s1-188	C	A	CTTAAATGGTGCTGGTTGCT GTATT	AGTGATATCTTAGTGGG TCGAGGAAA	AAACATGTAC GACCTGTC	TGTAAACATG TACTACCTGT C	0,0	omy07	564287 76
Omy_ar p-630	G	A	CTGCACAACCTGTTTCCTGCT ATT	ACCAAGTGTCCTGTAA GCC	CCGCTCCGTCT GCT	CCGCTCTGTC TGCT	0,0	omy07	572397 51
Omy_m yclarp40 4-111	T	G	GCTGTGGTGCTCATGGGTAA A	CCAGGGCAGGGTTGTTC TC	CAAAGCCATA CGTGGCC	AAGCCATCCG TGGCC	0,0	omy07	572399 57
Omy_R AD2976 -26	G	A	AGGACTGTGATCCTCTCAGC T	AGCTCTGCTGAAACATC AGTCT	CAGCTGGGTT GAGA	CAGCTGGATT GAGA	0,0	omy07	583371 26
Omy_97 077-73	T	A	GTGTAAACAAAATGACTCTG GGATTGAG	AGAAGTGGCAATGGTGT GAAGTAT	TGGTGCAATA GAAATA	CATGGTGCAA TAGTAATA	0,0	omy07	618754 65
Omy_R AD7384 -50	T	C	GACACGCCCTCAGCCAG	CTGGTACCTTCCTGCTG TGG	GCCTCTGGCA G	GCCTCCGGCA G	0,0	omy07	652417 84
OMS00 057	T	G	GAGAAAGGGAGCATGAGAC AGA	GTTGGGCTCCGGTACGA T	CTCCACAGAA CCTTG	CTCCACAGCA CCTTG	0,0	omy07	679081 35
Omy_10 5105- 448	C	T	CAATTTGCAAGCAGGGAAAG GTTAT	GTGATGGGCTGCAATTG CTT	AAGGAGAATG CATAATC	TGAAAGGAG AATACATAAT C	0,0	omy07	683035 84
Omy_cd 28-130	T	C	CACAACTCCACAGAGACAGT GA	GAGGACAAAACCTGACC GTATGGT	CCTGTTTCATTC ACCC	CTGTTTCGTTT ACCC	0,2,7	omy07	190XX XXX

OMS00 132	A	T	GTTTATGACTCCATTGCCGA AATGATT	ACGCGACCTGCAATTCA TCAATA	CAGCAGTCCT CTGTGTGG	AGCAGTCCTC AGTGTGG	0,0	omy08	945135 0
Omy_hs p90BA- 193	C	T	GGAATCGATGACGACGAAGT GATC	TTCCTCCATGCGTGATG CA	CCTCCGCGCCT GC	CCTCCGCACC TGC	0.5,0	omy08	106673 61
OMS00 153	T	G	ACTTTGCACCATAGGCTTGA CAT	TGATAAGGATGATCAAA AAGCTGAAGTATGTA	ACAAAATGTA ATTTTCC	CAAAAATGTCA TTTTCC	0,0	omy08	163033 16
Omy_hs f2-146	A	-	CCAACAATTGCAGCCTCATC TTAAT	GGAGCAGAAAAAAGGAT TGGACCTT	ATAATCTACT A	ATAATCTAAC A	0,0	omy08	174703 64
OMS00 068	A	G	GCACTAACTGGACAACATTT TTAAGAATGA	GGCAGTTGAGCATTTTG GGATATT	AATATGCCTC CTTCGTCTC	TATGCCTCCT CCGTCTC	0,0	omy08	174880 38
Omy_sr p09-37	C	T	TAGTTGTATTAACCTCTTCTTT GAGTCTAGA	TCATTCCAGCTCCGTTCT CTTC	TTGTGCTATTG ACGCCACAG	TTGTGCTATT GACACCACAG	0,0	omy08	214268 23
OMS00 151	A	G	CTAACGTCTTCCCAATGATA TTTCACAAGATA	ACCGTGGAAATACAATT TTTTATGCCAAT	TCATGACCTTG ATAATC	ATGACCTCGA TAATC	0,0	omy08	239492 30
OMS00 179	A	C	GTCATAACAAAATCAGGGCT TTCCAA	TGGGAGATTTGGGCTGC TTTAAA	TGCCTCTTCTC TTTTCTCAT	CCTCTTCTCTT GTCTCAT	0,0	omy08	255399 30
Omy_R AD2428 7-74	A	G	ATTGTCTGTCTGCCGAGGTG	TGGCGACCTGTCACTAA TGC	GGTCACTACC TCCC	GGTCACTGCC TCCC	0,0	omy08	280357 34
Omy_12 0255- 332	A	T	GCTAGCTAACATTGAAGGGT GGAAT	GGCTACAGGGACTTTAC AATGGG	ACTATGCCAT GAAGTTA	ACTATGCCAA GAAGTTA	0,0	omy08	291028 20
Omy_R AD2823 6-38	T	C	GGCACACATCTGTCCCGTAG	GCACTAAGGTCTAGGAG CACG	ATCTGTCTTCG TGC	ATCTGTCCTC GTGC	0,0	omy08	403994 99
Omy_R AD8670 6-72	C	T	TTCCCTGTAACTGTCACGCC	CCACATCACACCCTGAC CTC	TACGTTTCATT TCT	TACGTTTTAT TTCT	0,0	omy08	404794 21
Omy_10 4519- 624	T	C	CGTGTGAGTTTGCGGTAAAG AC	TGACGAGTCCGTCTTAT CATCCT	CAGCAGGATA CATCCGACT	AGCAGGATAC GTCCGACT	0,0	omy08	427641 23
Omy_10 8007- 193	A	G	GTGAATACCACCCAGGCTTG T	GTCCCTTCCCCAGTTTC ACTTAATT	ATGTTTTCTCC CTACTTAAC	TTTTCTCCCC ACTTAAC	0,0	omy08	436719 91
Omy_co x2-335	T	G	AGCTGGGCTGTATTTGTCAA TACTT	CAGCCCGCCACTGTCT	CTTTAAAGAC AAAGACTTTA T	TTTAAAGACA AAGCCTTTAT	0,0	omy08	508226 29

Omy_R AD1890 3-48	A	G	GGGGATGAGTTCTTCGGTGG	CCACCAAATCCCCGAA GAA	AGGAGACACC A	AGGAGGCACC A	0,0	omy08	610392 08
Omy_nk ef-241	C	A	AGTGTCATTGATGTCGGCCT ATTTT	AAACGAATGTCCACCTC AGATGTT	CTTCTGTATCA TTTTTG	TCTTCTGTAT AATTTTTG	0,0	omy08	694270 37
Omy_97 865-196	A	G	TCCAGACTTCTGGTTTGTTC ATT	CCAGCCCCTATATTCAC AATTAAGTGT	ATTAATTAAC AAGCTC	ATTAATTGAC AAGCT	0,0	omy08	825133 44
Omy_rb m4b- 203	-	T	CTGAAATTTGATGAATGGAA GCTGCA	CGTATTCAAGTCGATAT ACAGTCACGAT	CACGTTATTAT GAAAAGGATG T	ACGTTATTAT GAAAAAGGA TGT	0,0	omy09	124851 17
Omy_11 2820-82	G	A	CCTTTCCTTTTGCATTTCTC TACTTATTTATTT	AAATGAACTCACGTTGA CCTCTGA	CGCCGCCAAG TTA	CGCCGCTAAG TTA	0,0	omy09	218612 64
Omy_R AD6595 9-69	G	A	ACATTTTGGTGTAAACAACC CTGT	GCTAGCGAAGACCCTGA AGG	TTTTGTCTGTT CTT	TTTTGTCATTC CTT	0,0	omy09	363384 72
OMS00 103	A	T	GAGATCACTGTAGGATTGGC TGTTT	CCTCAGAGCAGCTCACA ATGGCATC	CTCCACAGTA ATTTTTTTTT	CCACAGTAAT TATTTTTT	0,0	omy09	383356 92
OMS00 056	T	C	TCAGGAAGTAAACTGAAAAT TCCAATGTATGA	CCCCAACCATGCTTGTT ATTGAAC	TAGCTTGACC AAATAGCA	CTTGACCGAA TAGCA	0,0	omy09	437713 25
OMS00 175	T	C	TTGCGATATGGGACTGTATA CATTTATTCC	ACTACCTCCAGTTAAAA TAGTGTGGGAAA	ATCACTAGTTC AAATACAA	ATCACTAGTT CAGATACAA	0,0	omy09	503817 07
Omy_tlr 5-205	T	A	GAGCGTATCTGGTATGGTAA CAACA	CTCCAGCAGCTTTAGAG AGTTTACA	CAGTAATATTT CAGTGCCCG	CAGTAATATT TCTGTGCCCCG	0,0	omy09	605104 83
Omy_11 4315- 438	T	G	CCTCACCGATCTAGTCAACT TCATC	AGGAGGCTGAGGGAGA TTCTAG	TTATGGGCTTA AGGGTC	TTATGGGCTT ACGGGTC	0,0	omy09	645645 08
Omy_gl uR-79	C	T	GACTGTCTATAGCTATTCTTC TCAAACGT	AGAAACTACCATTGTGA TTAACAGATAGAAAATA CAT	CAAGTATTTTG CGTAGGAAT	CAAGTATTTT GCATAGGAAT	0,0	omy10	750822 1
Omy_R AD3213 9-58	G	A	GCAGGAAACAGGTACAAAG GA	TGGCTTCTTCCTTGCTGA GC	TCGACATGAC CTGA	TCGACATAAC CTGA	0,1	omy10	107738 03
Omy_10 9894- 185	T	C	GGGAGGAATTGGAATGACA GATTAAC	CGGTGTCATTATGGTTG TCATTGTG	CTCCCTGATCC CCC	CTCCCTGGTC CCCC	0,0.5	omy10	108522 82
Omy_R AD6640 2-36	T	C	GGTGTGATACCTCAGAGCTC TG	CGTCTCCGGATCGTTCA GAG	AACCACTTCTC TG	AACCACCTCT CTG	0,2	omy10	303720 84

M09AA E.082	T	G	CTATGTGCAGTGCCCTTCTCA	GGCTTACAAGTATGCAT GACTAGCT	AGGTTGTTTTA CAAATTTAA	AGGTTGTTTT ACACATTTAA	0,0	omy10	328727 04
OMS00 095	A	T	CTCCAATGGCTGTCAACAAT TAAATATAAGAC	GTGTGCTGGTCTCTTCTT TTATTCTCA	AGGCAACTAT ATATTTTTTT	AGGCAACTAT ATATATTTTT	0,7,0	omy10	387046 54
Omy_R AD2608 0-69	G	A	TGTGGGACAGCACATACTCC	CCAGGACACCAGTGGA GAAG	ATTAGTAGCA TCATCGAG	ATTAGTAACA TCATCGAG	0,0	omy10	404866 18
Omy_R AD1073 3-10	A	G	TATAGACCCCCTGCCAGTCA	ACAGAGAAACCCCCGTC ATT	AGGGTGAAGA ACTG	AGGGTGAGG AACTG	0,0	omy10	414117 54
Omy_ftz f1-217	A	T	ACAGGGATGGGCAACTTTGT T	GGATGACCCACGTGACA CT	TCATGACGAG TTCTGATTT	TGACGAGTTC AGATTT	0,0	omy10	440609 19
Omy_p5 3-262	T	A	CCCCAACATCCAGTATACAG TTTCA	CCCAAATTGGCAATTTT AATAGGATTGAGA	CAAGTAGTAT GGAGCTCTAT	AAGTAGTATG GTGCTCTAT	0,0	omy10	461908 22
Omy_tlr 3-377	C	T	GTCGCTCCGGGTGCTT	GGCCCAAACACTTCCTT CCT	CGTGATTAGG TTCTTC	CGTGATTAGA TTCTTC	0,0	omy10	564470 78
OMS00 106	T	G	CGTGTAGCATTCTTGAGGAA GCTT	TTTCCAACAGATGCCAG AATCCT	TCTGATGGAA ACTTTC	TGATGGCAAC TTTC	0,0	omy10	598556 97
OMS00 030	T	G	CCTCGTGACTACAGAGCTAT ACAAC	GATCTGATCGGTCGGGA GAGA	ATGAGGGTCC CTATACAGG	ATGAGGGTCC CTCTACAGG	0,0	omy10	604014 05
Omy_U T16_2- 173	C	T	ATTGACTCATTATCACCTTAG TTGTAGCTTCA	GCAGCTACTTGCTGTAT CACATGTTTGT	ACAGTCAACA AGGGACTTAA	ACAGTCAATA AGGGACTTAA	0,0	omy10	614962 64
Omy_R AD3684 8-7	G	A	CGAGGACGTTCATAGGGAGC	TCGATAAGTCCACCAGC TGG	TGCAGGGACA CCACCCT	TGCAGGAACA CCACCCT	0,5,0	omy10	636507 47
Omy_R AD5374 -56	A	C	GCTGTTACCGTGTGATGTTG A	AGAGTTCTGGCCTCTCC CTC	AGAGGGAAAAG AGAG	AGAGGGACA GAGAG	0,0	omy11	618693 7
Omy_g1 2-82	T	C	GATCAATTCGATCGCTCATG AAACTT	CTTCTCTCGTTCTCATTG TGTCTCA	CAAACCTCTCA GGATTAG	AAACTCTCGG GATTAG	0,0	omy11	147648 07
Omy_B AMBI4. 238	T	C	CATGATGAGGAGGACCAAG ATGAG	AGGTGTGGTTCAGGGCA G	CACCGCAATC ACCG	ACCGCGATCA CCG	0,0	omy11	244110 85
Omy_M YC_2	T	C	CGGTTGCAGAACTCTCATGT TTG	CACGCCATGTCTTAACT TGCATTA	CATAGACTTTT TGACCTTAT	CATAGACTTT TTGGCCTTAT	0,0	omy11	338412 60

Omy_R AD7850 2-57	T	G	GAGAGGCATCCTGTCTAGGG	ACCATGCTCTTTCTGTG GGTGT	GGAAATATCA CACA	GGAAATAGCA CACA	0,0	omy11	370317 57
Omy_B AC- F5.284	C	T	CCTCATTTACTGTAGGACCA TGCA	ACAACGCCAACAACCTTT CTCTTG	CAGTAGGGCG GCAAG	ACAGTAGGAC GGCAAG	0,0	omy11	427893 02
Omy_co x1-221	T	A	CACTGAACTGTAAGCCATTG TGATT	GCAACATGGGAATGATT CATAAATGCA	CGGTAAGACC ATTAAAA	CGGTAAGACC ATTTAAA	0,0	omy11	470523 10
OMS00 120	A	G	GGCAGAAGAGGAGAGAGAT ATGATTG	CCTCAAATACCTCTGAC ATTGAAGGTT	C[GA]CCCAC AAAAC	C[GA]CCCACC AAAAC	0,0	omy11	510131 77
Omy_R AD4279 3-59	T	C	CACGGCTAGTGGCATGTACC	CCACACCTGCATCAGTC TGT	CAGAGAATGC CAACAGA	CAGAGAACGC CAACAGA	0,0	omy11	514085 22
Omy_10 2867- 443	T	G	CATTTGTTTAATTTGATTTGG CACAACCTCA	CCCTAGTTCTGTAACAC AAGACGTA	TTTGGGTACAT AATTTTT	TGGGTACATC ATTTTT	0,0	omy11	559343 32
Omy_ox ct-85	A	T	CGTCACTGAAACATTACTGT AACATCCA	CATCATCACGCTGTTGG TTTCTTAA	CATCGCTTATT TATGC	CATCGCTAAT TTATGC	0,0	omy11	684058 94
Omy_11 7540- 259	T	G	GGCAGGTTAACACAGTCATC TACTATAAA	CAGCATGTTGCTTTAAT CCTTCACA	TGTCACTTCAA AGTTTG	TGTCACTTCA ACGTTTG	0,0	omy12	507937 1
Omy_11 0201- 359	T	G	GGTAAGGCCTGTCTGACTAT TTTGA	AGAGGTCAATGGATGCC AGTTT	TTTGGCTATTG AAATTATACA TT	TTGGCTATTG AAATTCTACA TT	0,0	omy12	287279 52
OMS00 074	T	G	CCTGTTTATTCATCTAAACCA GTTCTTTAAAAT	AACTTAATTTAGCAAAC AAATGTCTGAACAGAA	TGAAACAAAA CAAATGTTCC	AAACAAAAC ACATGTTCC	0,0	omy12	311047 27
Omy_hs c715-80	C	A	CCGGTCTACCCTATAGCTGTT G	AGTCAGTCAATTAGTGG TTTGAAATACTATCA	AACTGTATTTG GGAAAAT	ATAAACTGTA TTTGTGAAAA T	0,0	omy12	401363 09
Omy_R AD2669 1-36	A	G	TGCAGGAAACCGTCAATCTA CA	CAGGAATTAATTGTATG GCCGGA	TCTCCTAACA GAAC	TCTCCTAGCA GAAC	0,0	omy12	472849 87
Omy_R AD3840 6-19	T	A	CTGCAGGGGTATTAGGAGGC	AATGAGTTGTGGCGGTG AGT	AGGCTTTATAT GGCC	AGGCATTATA TGGCC	0,0	omy12	481811 30
OMS00 077	C	G	AATACCATCTTGAGCTCATT AGTAATTATTCAA	CCAGACTTTACACACTC TTGACTGA	TTCCGGTGGT GAAGTT	CCGGTGCTGA AGTT	0,0	omy12	488619 50

Omy_R AD1919-22	A	G	CAGGTCACAGACACACAGGG	CTACACCACCCACGTTCTG	CAGGGAGGAGG	CAGGGGGGAGG	0,0	omy12	53383757
Omy_ndk-152	A	G	AAGAATTGAGGGATAAAAA CAAAATAATATATAAACATGA	CAAACCTACATTCATTA AAGTCCAGTTTTGT	ACCCACTTTCAAAAC	ACCCACTCTCAAAAC	0,0	omy12	56277959
Omy_R AD76570-62	T	G	GCAGGTAGGTAGGAAGGAAAGC	TCTGACTGGTATTGAAA GGACCA	AGAGGTGTTCTGGT	AGAGGTGGTCTGGT	0,0	omy12	56297752
Omy_gh-475	C	T	AAGTTACCAGAATTTTGCAA ACTCAACT	CCATATTTTGAGGTGTA GCTTTACCCT	CTGAAACTCATGGTATACA	CTGAAACTCATGGTATACA	0,0	omy12	62308122
Omy_R AD33122-47	G	C	CAGGCTTTGTGGACATGTGC	GTGCTCTATCTTGCTCTTGGC	CCACAGGGTGGTGC	CCACAGGCTGTGC	0,0.6	omy12	66078752
Omy_hsp70aPro-329	A	G	TGCGTATTATTGTTTTTCAAG GACTTTCAAA	TGAATATTTTCAAATAC ATGCCAATTCTTTCCAA	ACATTCCAATATTCAACTAT	CATTCCAATATCCAACTAT	0,0	omy12	66828836
OMS00053	T	C	GGAGCCAGGTCAAGGTGATC	GGATGTCTGGTGTGGCTGTAAA	ATTTATATGTATCAATCA	ATTTATACGTATCAATCA	0,0	omy12	66917868
OMS00112	A	T	TGGCAGCAAAAAGGGATGCA	TCCTGAGCAACCAGTCAACATT	CCGGTTTCAAGTTTACTTGT	CGGTTTCAAGTATACTTGT	0,0	omy12	68382081
Omy_11 1666-301	T	A	GGGTGAAAAGAGTGGGACATTTACA	GTCAATTTCAAGGCACCAGACAAT	AGTATAACACAGTAAGACAA T	AGTATAACACAGTTAGACAA T	0,0	omy12	70990844
OMS00149	T	G	GGCATCATTGTTCTTGCTCTGTTTA	CCTGGGAGGGTTTATATCGGAGTAT	GCTAAATGCACAG	GCTAAAGGCACAG	0,0	omy12	6589XXX
Omy_R AD66834-17	C	T	CTCCTGCAGGTCATCTCTGG	CTGTCTTGCTCAATGCCCTG	TCTGGCTGACACCTTTA	TCTGGTTGACACCTTTA	0,0	omy13	10112620
Omy_11 8175-396	T	A	AGGCTTCACACACACATGCA	GACGCGCAACCTCTAGATTATACTT	CTCTTGACAGACATACCCGTA	CTCTTGACAGCATTCCCGTA	0,0	omy13	20282478
Omy_12 9870-756	C	T	TCGTTATTTTGCCCTCGCGGTA	TCCCATGAAGATGTATACATGTTTGTGA	ACAGGTATTTCTGAAATG	CAGGTATTTATGAAATG	0,0	omy13	22915161
Omy_11 3490-159	C	T	CATAGTACATTTACAGATAATGTTTTAAAGTGCATGT	CGAGATACCAAAATGCCACAGTTACAT	CATCTGTTTTGTGTTAGC	CATCTGTTTTAGTTTAGC	0,0	omy13	26494831

Omy_nach-200	A	T	CTCATGAAAAACGGGAGAGCAAAG	CAGCGGCTCTTCAGTAGTCT	AACTGACAGAGTCACAAC	CTGACAGAGACACAAC	0,0	omy13	30001796
OMS00180	T	G	GCGCCGAATGGCATTAGG	CACATTGCTGTCGTTTAGTTGACT	CTAAAAGTGCATTAAGCC	CTAAAAGTGCCTTAAGCC	0,0	omy13	32462775
Omy_110064-419	T	G	GTGCAAGGGACCTAGCTAATCC	TCTGAACTGACACTGAA GAACAAAGAA	ACGTTAGCTTTTAATTTTC	AACGTTAGCTTTTCATTTTC	0,0	omy13	36272850
Omy_g1-103	T	C	AGTCGTGACAATGAGAAACAGTGTT	CTCAGCAAAAAAGAAACGTCCCTTT	CCTTTTACAATGAAGATC	CTTTTACAGTGAAGATC	0,0	omy13	39287232
Omy_IL6-320	C	T	CGACTGATCTCCTGCAGACATG	CTTGTTCCCTCGTTGTCTTCCTTCTA	CTATAGGAGAGAGGACAACA	ATAGGAGAGAAGACAACA	0,0	omy14	7102407
OMS00089	A	G	GCACCATTTGAATAAAAAATCTGCTTTGT	GCAACCCAATTCAATATTAAGCACATGAT	ATGAATCCCAATAAGAAC	AATCCCAAACAAGAAC	0,0	omy14	13150534
Omy_hsf1b-241	A	-	AGCCCGAACTATCCTAAAGCATTTT	AAATCAATAGCTCAGAG AATAATGAACACCA	CAGTGTTTTGT TTTTGTGTCATT	AGTGTTTTGT TTTTGTGTCATT	0,0	omy14	14516742
Omy_116733-349	C	T	GAAATGGACATGCCTACAAATTGCT	GATGTGATCAGTTTAGGCAAGGC	AGAGAATCTGATAGTATTTTC	AGAGAATCTGATAATATTTTC	0,0	omy14	18498042
Omy_ntl-27	G	A	GGTGTGTTACTGTAGTTGTGTCCTT	TGTGTAGCTAGTGATCC TGATTGTCT	CAGACAAGAGTACCCCAAGAC	CAGACAAGAGTACTCCAAGAC	0,0	omy14	22071034
Omy_txnip-343	T	C	CCTTCAAACCTAACGCATCATAGACATG	GGTCACTTGGCTAATCC CCTTAT	AACTGAAGAGATCTG	AACTGAAGGGATCTG	0,0	omy14	24435825
Omy_UBA3b	A	T	GCCACTCAATGCATGTGTTTTCTAG	CAGCTAGCTTAAGTGGGATGCAA	TGGAGATAACGCTAACTATT	AGATAACGCATAACTATT	0,0	omy14	28552649
Omy_RAD55404-54	C	T	GCAGGGTGTCCACTACAGAC	AGGAGTCCTGAGAGTTGGC	ATTGTTTCTGAGG	ATTGTTTTTGAAGG	0,0	omy14	43876869
Omy_RAD12566-14	C	T	GTGGACATTCCTGCAGGGAT	TCCCACAAATATTTTCATACGCACA	ATGTAAACAAATTG	ATGTAAATAAATTG	0,0	omy14	44125336
Omy_RAD3926-22	T	C	CGTTCCTGCAGGCTTTTCAC	TTGGCACAGAGAGTACGCAG	TTCACTTTTCCCTG	TTCACTTCTCCTG	0,0	omy14	45345547
Omy_mcsf-268	T	C	CCAGCATTCGTTCCCATTTCC	CTTTTAATGTAGATTATATTCTTCTGTAGCCACTATGG	AAATAATAGATAAA[CT]CCT	AAATAACAGATAAA[CT]CCT	0,0	omy14	48758470

Omy_10 1554-306	T	C	GCCTGTATTTCTCCTGTATGT GCAT	TCAACTTTTGCAAACCTT TTTTATTCTTTGTCAATT	TGCTTCTCACA TTTTTA	TGCTTCTCAC GTTTTTA	0,0	omy14	536601 17
Omy_R AD4744 4-53	C	T	GTCGTCTGGAGGAGCTGAAG	GGGTGACGTTTTTCCTTC AGC	GGCGAGCTTG GCCCCAAA	GGCGAGTTTG GCCCCAAA	0,0	omy14	560519 17
Omy_R AD1763 2-23	C	T	AAGCTCCTGCAGGTCATCTC	TCTGTGAACTGTCTTCT GCAAGT	CATGTGAGAC CTTTGCA	CATGTGAGAT CTTTGCA	0,0	omy14	560744 85
OMS00 072	A	G	GTGGGAGAGCTCGTCTATGG	ACAACAGGTCATTGGAT GTGATCAG	TAGAAGGTCC ATGTATCTC	AAGGTCCATG CATCTC	0,0	omy14	599717 80
Omy_re dd1-410	C	T	GTACTCCCACTAACATACAG TAGACTCA	GGCACCATTGTGTTTTA GGATGTAG	AAAATATCCT GCAAGGAAT	AATATCCTGC AAGAAAT	0,0	omy14	690441 21
Omy_nx t2-273	C	T	CTTTAGAAAAGCCAAGGTAT ATTTTAACATACTTCT	CTGCTGCCCTCTAATGG TAAGATAG	AAGGCAC	AAGGCAT	0,0	omy14	719697 84
Omy_11 0689-148	A	C	GTGTGTGGCAGAGAACTAAC TGAT	GGTTAAGACATTAACAT AACACTGGACTCT	CAAATGAACA CATTATTTATC	ATGAACACAT GATTTATC	0,0	omy14	7235X XXX
Omy_R AD8802 8-7	G	A	TAGCCCAGTTCGGTTCCAAC	AGTGTCTTTGGTGCGTC CTC	TGCAGGGGCT GG	TGCAGGAGCT GG	0,0	omy15	567666 4
Omy_O go4-212	T	C	TGAAAGGTTTTATGCAGGTT ATTTTCT	GTGTGTGTTAAATAAGC ATTTGATGA	CATTTGATGA GACATCTT	ATTTGATGAG GCATCTT	0,0	omy15	125958 06
Omy_hu s1-52	G	A	CTTGCCGGAGGGTAGCT	CCACAACCTTCTCAAATG AATGGAATGT	CCCATCCCTCC TCCTGG	CCCATCCCTT CTCCTGG	0,0.5	omy15	130422 22
Omy_11 1084-526	A	C	CACCACACCAAGCAACTATT TCATT	ACCCAACACTGTCCCA TTTTTCAT	CCAGTGAAAT TTATTTTT	CAGTGAAATG TATTTTT	0,0	omy15	173008 50
Omy_11 1383-51	C	T	CACGCGCAATCTCTCGTTTTA C	TCTTTAGGCAACAAGCG TGTCAC	AGCAAGCGCA CT[AG]GGT	AGCAAGTGCA CT[AG]GGT	0,0	omy15	212397 68
Omy_ca rban1-264	G	A	GCAAAGCCTCATCTTCAATC ATTTGT	GCAAAACACAAGTCAG GAATCACTTA	CATTAATATTG CTAATAACAC CAAG	ATTAATATTG CTAATAACAC TAAG	0,0	omy15	214540 64
Omy_96 222-125	T	C	GTAAGGAACTAATTGGCGCA ACATT	CAGTTTGTCTAACACCC AGGCATAT	AACTACAAC GTAGCTAATT	CAACTGTGGC TAATT	0,0	omy15	240411 11
Omy_R AD2357 7-43	T	C	AATAGGAACCAAGCCCCAGC	CAGAGCCTGAACCCATG GAG	TCTGGCTCTGT CGGTCT	TCTGGCTCCG TCGGTCT	0,0	omy15	269722 88

Omy_R AD5281 2-28	C	G	AGGAGTCCTGTCCCATGTCA	GCTTAAGGCTGTGGTAT GTGG	CAACCTC[TC] ATTCCACAT	CAACCTG[TC] ATTCCACAT	0,0	omy15	286237 29
OMS00 061	T	C	AAGTGGAGGCTGACCTGTTG	GCTGATGGCACCTGACA GTTAATT	CATTGCCATTT ACAGACTT	TGCCATTTGC AGACTT	0,0	omy15	312319 75
Omy_ni ps-299	T	-	GACAGGATAGGAACGGTTTC TCAAT	ATCAGAAGTTTAATTCA ATATGTACACGATCCT	CTGGATTTCAC ATGTAATAC	CTGGATTTC CGTAATAC	0,0	omy15	397887 84
Omy_L DHB- 1_i2	C	T	ACGCACACTTATCCTTGACA ATGTT	ACTGTGACAACAAATTC GGTGACA	ATGGGCAGTC ATTCA	TGGGCAATCA TTCA	0,0	omy15	440944 40
OMS00 143	T	C	GGAGGCACGCCCCAAA	TTTGTAAAAATAGAGCC CTTAGTGGGTTT	CCTGATCCAG AATCTAGA	CCTGATCCAG AGTCTAGA	0,0	omy15	592330 15
Omy_97 954-618	C	T	GCTCTGCTTCCTCGGCAAAT A	CACAATTGGTTTTTGCA CAAAAGTAAAGTATT	CAACGCTTAC CGGTGTGT	CAACGCTTAC CAGTGTGT	0,0	omy16	114334 35
Omy_11 7370- 400	A	G	TGCAAACACAGAGGAAAGG GATTT	GGCTTATTTGTTCCGTA CTTGCAAT	CAACTCCAAT GAATTAA	AACTCCAACG AATTAA	0,0	omy16	148539 62
OMS00 062	T	C	ACCCTGGGAAGGCTACTGTA C	TGAACAGAGATCTGGAG AGTTGGAT	TTGACCAGCA GATGGTGTA	ACCAGCAGGT GGTGTA	0,0	omy16	148580 81
Omy_R AD8812 2-32	G	A	TCAGTGGATGGAGTGCCCT	GGTCTTTGGCCTTGTTG CTG	GCTGTGGAGA TCAT[CT]CG	GCTGTGGAAA TCAT[CT]CG	0,0	omy16	163340 74
Omy_R AD4064 1-58	T	C	GGCAAACCTGGCTTGTGAGTG	AAGGCTCTGCTTCTGCT TGA	AGTGATATCA AGTG	AGTGATACCA AGTG	0,0	omy16	178582 78
OMS00 041	G	C	GATTCTGTTCCATCCTCTTTC TGTCA	AAACATAAAAAAGGGC ATGAAGGTGTC	CCACTCTATGC CTGCCCT	CACTCTATGC GTGCCCT	0,0	omy16	182791 74
OMS00 119	A	T	AGCGGCAGTTGTGTTAATGA GA	CTTCCTAAAGCCTGACA GTCTGT	CCACACAGCT GCCTGT	CACACAGCAG CCTGT	0,0	omy16	275060 18
Omy_R AD116- 59	T	C	GGAAGAAGTGAGAGCCCTG G	CTGTAGTCCACGATCCG CTC	CCACAATGTC AAC	CCACAACGTC AAC	0,0	omy16	330748 66
Omy_R AD1784 9-16	G	C	GACTCCACAGCCTACATGGG	CCGTTAATGCCAGGGGA GTC	AGACGGACTC CCC	AGACCGACTC CCC	1,1,0	omy16	441164 56
M09AA J.163	G	A	TCCCATGGCCCTTACTCTATC AA	TTGAGGTGTATGTTGAA AAGTAACTT	AACAAAGTGA AAGTGCCT	CAAAGTGAAA GTGTCTT	0,0	omy16	448434 40

OMS00 018	T	G	AGAGTACATGTGTGGCTGCA A	GTCATAAATCAACACAA TTATCTTCTTCACAGAA	AACCACATAA TTAATAATTC	CCACATAATT CATAATTC	0,0	omy16	464324 62
OMS00 134	A	G	GAAACTGAAATGATCCCATC GTGTT	GCTAGCATAACAGCATT GCCATAT	TCTATAGCTGC AGTATATTA	TAGCTGCAGC ATATTA	0,0	omy16	477436 37
Omy_R AD3140 8-67	T	C	CAACCCTGCAGGCTACAGAA	TGGAGTGCCAACAAAA GAAGC	ACAGAATGCA GAAA	ACAGAACGCA GAAA	0,0	omy16	483410 10
Omy_R AD7931 4-58	C	T	CACACTGACTCATCCCTCGC	GAGTGTCTTACCGAGCT GCC	AGACCTTGTC	AGACTTTGTC	0,0	omy16	627284 32
Omy_R AD4013 2-55	A	C	TGCAGGGCCTGTATATTGCT	TCAAAGGACTGGGGAG AGGA	TCTGTGCAGTC CTC	TCTGTGCCGT CCTC	0,0	omy16	627847 79
OMS00 006	T	C	TCCACGTAGGACATAGTTTG AGCTA	TGTGGTGTGTCATGTTTGC CCTAC	CACTTACAAA TACAAAATT	CTTACAAATG CAAAATT	0,0	omy16	632479 44
Omy_an p-17	C	A	GGTAATGCCACATGCGGTAA ATT	GGCGAAATCTGAAAATG TGCTGTTA	CTCTCATTGGT ATAGTAACC	CTCATTGGTA TATTAACC	0,0	omy16	3072X XXX
Omy_10 3705- 558	T	C	CTCCAATCGCAAATACCCAG ACT	CGCAGGAGACGGATGC C	AGACTTACCC AGAGTGAGAG	ACTTACCCAG GGTGAGAG	0,0	omy17	706598 6
Omy_11 6938- 264	A	G	GTTTCATTTCATGTTGAAGTGC GACAT	CTCTGCATGCTCCCATC CT	CCTTGTCTCAA TTTTTCCTCT	CTTGTCTCAA TTTCTCCTCT	0,0	omy17	715446 3
OMS00 128	T	G	ATGAAAGAACTCCCAGACAC GTATTTT	ACATTTTAACACAGTAA CACTAATACACACCA	ACTCTCAGAA TTAATTATG	CACTCTCAGA ATTCATTATG	0,0	omy17	144885 87
Omy_10 1832- 195	A	C	TGGCTCTGGACCTGTTGAGA	CGTCACAGCTATTTTAG GCGTAGT	TGTAGTCTTTC AGAGTAGTAT G	TAGTCTTTCA GAGGAGTATG	0,0	omy17	170156 58
Omy_R AD4510 4-18	A	G	TGGTGCTTCAGTGCTGTCAA	AGAGTGAAAACGTGTGTG CGG	CAAGACACCG CACACAG	CAAGACGCCG CACACAG	0,0	omy17	206937 54
Omy_10 1993- 189	A	T	ACAAAACACAGTGGAATTAC AATTAACGTT	GGAAGTTAAATTTTCGCT TCGTCAGAA	CTTGATTGCA GCTTGTCAA	TGATTTGCAG CATGTCAA	0,0	omy17	214912 90
Omy_R AD2212 3-69	T	C	TGGGAAAGCATAGGAGGGG A	TGTGTGCCTGTCTTATA GCCC	CCAAAGATGT CAGA	CCAAAGACGT CAGA	0,0	omy17	237569 20

Omy_11 4976- 223	T	G	GACAAACAGCACTTCATTGC AGTAA	GTTGCTCCAGCACCAGG T	ACCGATGGAA CAATC	CCGATGGCAC AATC	0,0	omy17	414629 73
Omy_u0 9-56.119	T	C	CCAAGGTGGACCCACCAG	GCTGAGTTTATAGGTCA GTCATTATACATATTGA	AGTGAGCTGA AACAGAGCA	TGAGCTGAAG CAGAGCA	0,0	omy17	416909 56
Omy_ca 050-64	T	G	GTCATACAGAACTGTTTTGTT GTGTCAA	ACCTTGAATTGGTTCCT AATGCTATTGT	CAGTTTGAAG AATATACTC	CAGTTTGAAG ACTATACTC	0,0	omy17	486517 93
Omy_R AD4369 4-41	A	C	CCCCTCTCCCTGGCTAGAAT	TCAGGGGGTGTGCTTTT CC	AGGGAAGAGC GGAG	AGGGAAGCG CGGAG	0,1.1	omy17	577285 49
Omy_R AD4667 2-27	C	G	TGCAGGAGGTCTTTTTCCTTG T	AACACATTCTTATTTGC AATGATGG	GTGGTAGCCC ATCA	GTGGTAGGCC ATCA	0,0	omy17	582563 53
Omy_R AD5821 3-70	A	T	CCTGATGGGTGCTCTTCTCTC	AAACAGCATCATTATCC ATAGTGTT	TTTTTT[TA]AA AATATACT	TTTTTT[AT]TA AATATACT	0,0	omy17	582662 27
Omy_U 11_2b- 154	T	C	GGGAAGCAGAAAACTGGA AGTT	CCCTCTGTGGGCTTGAT ATTCA	AATGATACTTT TCAGATTGTA AC	TGATACTTTT CAGGTTGTAA C	0,0	omy17	594666 96
Ocl_gsh px-357	T	G	GAGATCCTGAGGTCCCTGAA GTAT	AAGTGGAAATTTGGGCT CAAAGC	ATCCGTCCAG GAAATG	TCCGTCCCGG AAATG	0,0	omy17	646977 03
Omy_G HSR- 121	T	C	CTGTGTATAAGTTTATACAG TCAGCACAGT	TTCAGAGAGAGAAATG GCAGAAAGG	CCTAATAACC ATGATAACAG C	AATAACCATG GTAACAGC	0,0	omy18	116628 01
Omy_R AD5995 0-44	G	A	GGAGCTCATATCGCCGATGG	GAACTCTGTCACCCTGC CC	GGAGGGGAAG GG	GAAGGGGAA GGG	0,0	omy18	172929 64
Omy_R AD4246 5-32	G	T	GTGGATCTTGGA CTCCAGGC	TAGACATCGGCCCTCAC AGA	CCAGGCTGGA AGAA	CCAGGCTTGA AGAA	0,0	omy18	250342 87
Omy_R AD7877 6-10	T	C	CACAGCTTCCTGCAGGGTAA	GCTTGCATGGTCTCGCT AGT	GGGTAATCCT GGCT	GGGTAACCCT GGCT	0,0	omy18	250343 26
Omy_sa st-264	G	A	GAAGTAGGGTTTGTGACCA TGTGA	TGGATTCCATTTTAGGC TGTAATACATCTT	CTAGCCAATG CGTCTAA	ATCTAGCCAA TGTGTCTAA	0,0	omy18	282520 83
Omy_R AD4361 2-42	T	C	GTGGAGAGGGATTTTGGGGG	TGACAGGACAAACACA AGCCA	AAATGTGTAT TTGTGTA	AAATGTGCAT TTGTGTA	0,0	omy18	291187 77

Omy_12 8996- 481	T	G	CTCATCCACACTGTACAGTA CAAGT	CATGCCTTCGTCTCATC AATAACAC	CAAACCTCAA CCAC	CAAACCGCAA CCAC	0,0	omy18	308021 01
OMS00 121	T	C	GGAAGGAGGTCCAGTGTGAG T	AAAATATGCAACACCAC TAAAACTGGAAAA	ACAGCGTGAT AAATT	CAGCGTGGTA AATT	0,0	omy18	342329 91
OMS00 127	T	G	CACCTTTCTCTCTCTCTCCAT CTCA	AGTGTGCTACACAACCT TAAAAAATATATATCTA TT	CACACACCCA AATGTA	ACACACCCCA ATGTA	0,0	omy18	362684 23
OMS00 118	T	G	GCTTATTTAGAGTGCATGCC AGATG	TGGAACCAATGGGACA GTCCTA	GCGGGGTGTG C[AG]CATT	GCGGGGGGTG C[AG]CATT	0,0	omy18	422122 99
Omy_R AD7210 -8	C	A	ACACCACACTCCACAAAGCA	GCGCCTTGGTCTCCTTC ATA	TGCAGGACTT GCTTTGT	TGCAGGAATT GCTTTGT	0,0	omy18	425586 54
Omy_R AD1934 0-24	A	G	GCAGGGAGCAGCATATACAT G	TGGGGTGATTTGAGTGA CAC	CATGGAAATA CATA	CATGGAGATA CATA	0,0	omy18	427018 41
Omy_R AD3209 -10	A	G	CGGAGGAGTTTGAGCAGTCT	CTTCTACCACCACCTCG CTG	CGGTATCCCT GGC	CGGTGTCCCT GGC	0,0	omy18	509849 53
M09AA D.076	T	C	ACTGTTACCACTCTCTCATCA ACCT	GGGTCCAGGAGGTTTTT AAACAACAT	CACCAACCAC TGGTGAA	CCAACCGCTG GTGAA	0,0	omy18	537175 12
Omy_II- 1b_028	T	C	ACTGTCTGGCTAGAGCACAT TG	ATCTTCTACCACCGCAC TGTTTTAA	CTGAGGCAAC TTTTGT	TGAGGCAGCT TTTGT	0,0	omy19	103295 30
OMS00 092	A	C	TCTCCAGGTGTATCTTGAGA AGGT	AGGGTTCACACAGGGA AGATATCAT	CAGCTGAGAA TAGGTTC	AGCTGAGAAG AGGTTC	0,0	omy19	125185 51
OMS00 017	A	G	ATTAAGTTCATACAAAAGTT CATCATAAATATTTTCCTT	GGAGAACAAAGGGAAA GAGAAGACA	TAGACCTCGG TGCTGTAG	CCTCGGCGCT GTAG	0,0	omy19	191115 91
OMS00 105	T	G	ACATTTGAAGTCAGTATGGG TGTTGAG	GAACCTCACCACAGTAC TAAATGCA	CTGCTATTCAA ATTGCT	CTGCTATTCA CATTGCT	0,0	omy19	202649 68
OMS00 133	A	G	GACCACTTCACTCATTCCTCC TTTT	TCCGGTTTACACACTTC ATGCA	CGCCTCCATCT TTGTGGT	CGCCTCCATC TCTGTGGT	0,0	omy19	237020 93
Omy_ra pd-167	G	T	CCCAACATGCTCTATTGCAG CTA	AGTTGCATAAGATGAAT CAATAAATTAAAAACAC AGAT	AAACAATCCC CCCCAAA	AAACAATCCC ACCCAAA	0,0	omy19	273624 24
Omy_12 8693- 455	T	C	GCCTGCAGGAGAAGGTAGA GTTA	GAAATGGAATGGACCCC AATCCT	CACTCAACTG ATACCC	CTCAGCTGAT ACCC	0,1.4	omy19	328900 59

Omy_R AD4911 1-35	T	C	GCAGGCTTAGCATTGCTGAC	GGAACCTGGGTGGGAG AATG	TTTCTTATATT TGA	TTTCTTACATT TGA	0,0	omy19	400501 11
Omy_R AD739- 59	C	G	ACGAGGCTTGTAATGCAGT	TGCCTTTATACCAATGT CTGCTG	GAGTTGGCTA TTTT	GAGTTGGGTA TTTT	0,0	omy19	529237 02
M09AA C.055	C	T	GTCTCCGACGTGTGGCT	TGGAACGAACCTGAGA ACATAAGG	ACCTCCACGC TGTCC	ACCTCCACAC TGTCC	0,0	omy19	538551 67
OMY10 11SNP	C	A	AGGCTGGTTTGGGATTCACT G	CGCCAAACACTAACTCT CTGTCT	CTTTACCTCGA AGACAAT	ACTTTACCTC TAAGACAAT	0,0,6	omy19	544462 25
Omy_B AMBI2. 312	G	T	CGAGCTCATGTCCGAAACTC AT	TTTGACAGCCTCAACTT CTAGGG	CCGAAAGTTC AACTTT	CCGAAAGTTA AACTTT	2,1,0	omy19	611XX XX
Omy_R AD2970 0-18	C	A	AATGGAATTGGCCCCAACCC	TCTCCATTGTGTGTAAT CATGGT	ACAATTCAAA TGATTTA	ACAATTAAAA TGATTTA	0,0	omy20	167319 6
OMS00 039	A	G	GTCAGTACTGTGTGTGTCTGT GT	CCATCTACATTGTCAGC AGTGTGA	GTACGTGTCTC TGACC	GTGCGTGTCT CTGACC	0,0	omy20	480049 5
OMS00 114	T	G	GGATGATGCTGTGAGTCGAG AAG	ACCTTCGCCACCCATGT TTTATT	AAACGTTTCA CATGCACC	AAACGTTTCA CCTGCACC	0,0	omy21	110354 95
Omy_99 300-202	T	A	CAGTTTGACCCGATGGTGTG A	GATTATGGCGTGGCCTT TTGG	TCAGGCATGA GAGAAA	ATCAGGCATG TGAGAAA	0,0	omy21	146970 73
Omy_ci n-172	C	T	CGCATGGGACAGGTGTGT	GAGAAAGCCTGTAGAA CCATGTCT	CGCTCACCCT GGTTAC	CGCTCACCCT GGTTAC	0,0	omy21	216026 18
Omy_va tf-406	T	C	TTGCTTCATTTTGTGATAACC TTGGG	TGCATGCTCTGACAAAT GTTACACT	ATGACTATCC ACA	ATGACTGTCC ACA	0,0	omy21	238062 20
Omy_L DHB- 2_e5	T	C	TGCTAGGTGAGTCAGAGGTA CATATT	GACTGGAAGGCCACCCA TAAG	TTTACCTGTCA AC	CCTGTGCGAC	0,0	omy21	241299 07
Omy_L DHB- 2_i6	G	T	TCCTCGCCAATACCATACAT GTC	AGAGTGAAGCTAACAC ACACATTTCT	CTGTGTTTTGC TTCCCCA	CTGTGTTTTG ATTCCCCA	0,0,5	omy21	241305 59
Omy_10 4569- 114	A	C	CCGAGGCCGACGTGATC	GCGCCTCGCTCATCATC A	CGCCACTCCG ACGCC	CCACGCCGAC GCC	0,5,0	omy21	321785 17
Omy_zg 57-91	C	A	CACTCATACACTCACTCACA AAGGA	AGCAGATAAGCCTTGTG AGTGAATCTT	CACAGACTGC ACAGCC	CCACAGACTT CACAGCC	0,0	omy21	327667 40

Omy_R AD3500 5-13	C	T	TGGTCAAAGTTGAGGGTGGT	CAGGGCCCTGATTAACC ACT	CCAACTCCCG ACGG	CCAACTCTCG ACGG	0,0	omy21	371884 35
Omy_b1 -266	G	T	TCATGTGAACTTTAATTGACT AGGAAGTCG	GATATGAAAATATCTGA AGAGTTATATTTGGGAA ATTGAC	TCTATAAACA ACATTTTTC	TCTATAAACA AAATTTTTC	0,0	omy21	412557 73
Omy_10 04	A	T	GAGAATCGGAGCTAATCTTA GTTATTGTGA	CACTTTATTGAGCTACA TGGCAAATCTG	CATGTGATGTT TTTTTGC	ATGTGATGAT TTTTGC	0,0	omy21	419442 18
Omy_10 2505- 102	A	G	CTGCAAACCTGACATGGTAGC AAAA	TGCTTGCTTTTTTAAAAA CAATCTCCCA	AACAGGATGT TTTTGC	CAGGATGCTT TTGC	0,0	omy22	775932 8
OMS00 173	T	C	TGGAAGTAGCTACTTAACAG GAAATGG	AACACGTGTGCTTGTTT TGTCAA	CATTAGCTTGT GTATGAACT	ATTAGCTTGT GTGTGAACT	0,0	omy22	910589 1
Omy_Ot s249- 227	C	T	CTATCTATCTATCTATCTATC TATCTATCTATCTATCTACTT ACTGAGA	CCCCTAGATTAAACCTG TCCAGTCT	CCCTCTGAGA ACTAC	CCTCTGAAAA CTAC	0,0	omy22	189174 30
OMS00 058	A	G	GTGACATTTGGAGCCACTGC	GCTAGGAGACAGAGGG TGAAAG	CAACACTTTGT ACCCCTC	CACTTTGCAC CCCTC	0,0	omy22	199221 39
Omy_R AD9358 0-37	T	G	AGGCAGAGGAGGGTTGTTTG	TGCAGAAAGTCAAATCAC GAACA	AGTCACCTGG GATT	AGTCACCGGG GATT	0,0	omy22	242478 65
Omy_IL 17-185	G	A	CCACCACACTCTGCAGCTT	TTGACGGGAATCCGAGA CTTC	AAGAATCTCA CCTGCCCCAT	AAGAATCTCA CTTGCCCCAT	0,0	omy22	272260 63
Omy_R AD1033 59-45	C	T	GGAGAAGGATGTGCTCCCTG	ATTTGGAGGTGGAGGGT CCA	CCTGTAACGC ACAG	CCTGTAATGC ACAG	0.4,0	omy22	405024 75
Omy_R AD4848 -14	G	T	TGTCCCTCTTCTGCACGATG	AGTTGGTAGCTCACTCT CTGT	GAGACAAGGA CAGA	GAGACAATGA CAGA	0,0	omy23	549364 8
Omy_10 7806-34	C	T	TCTTTGTCCATGCACATTGAT ATT	AGCACATTTAGTTAGCA GTGATGGA	ATTGGATGTC AGTGTCATT	ATTGGATGTC AATGTCATT	0,0	omy23	100954 18
Omy_R AD4879 9-69	A	G	GCTGAGCCACCTACACACAG	GTCTAACACTCGCAGCA GGT	CATCCTAGAA TAGAAGT	CATCCTGGAA TAGAAGT	0,0	omy23	215810 80
Omy_18 7760- 385	A	T	CGGCTATTCTCGCGTAAAAG CT	AAATGCAACCAGAAAC GGAATGTC	TCCTTATCCAA AATTATTGTGC	CTTATCCAAA ATAATTGTGC	0,0	omy23	236597 37

OMS00 024	T	G	CACATACAACCATCACCCCTT CCTAA	AGCATTGAGCGAAATTA CCAAGAGT	AA[AC]CCCAA ATTTTAC	AA[CA]CCCAA TTTTAC	0,0	omy23	314183 39
Omy_R AD3651 -48	G	T	GAGTACAGTGCAGTGTGGGG	CCTTCCTCTTGCCACCAT CA	GTTGGGAGAA CTTT	GTTGGGATAA CTTT	0,0	omy23	320298 60
OMS00 048	T	C	GGAAGAGCTGGAGAACAAC GT	TGCAGTTGACAGAGGCT TTCTTT	CAGCTAAACT CAGCAAAA	AGCTAAACTC GGCAAAA	0,0	omy23	371256 68
Omy_e1 -147	G	T	GCACTGACTGTTACCAGGAA AGAG	GTACTGCAGTGTGAGG CTATATCA	CCATCCTGAA TCTGATTAA	CCATCCTGAA TATGATTAA	0,0	omy23	383133 38
Omy_10 9243- 222	A	C	ATGTGCACCTCTTAAATTGT AAGTAAAATGT	ACCCTATATTCAGTGGC AAGATTGC	TGTTTCATTAAA TTGACTTTTT	TTCATTAAAT GGACTTTTT	0,0	omy24	700968 7
OMS00 101	A	G	GCGTGTCTGTTGGGTCAGTTAA ATA	GTGCAATCCAACCTATT AGTAGATATGCT	CTCTAGTAGC CTTATAGAAA G	CTAGTAGCCT TACAGAAAG	0,0	omy24	104754 51
Omy_in os-97	C	A	GATGGACAGGGTCTCTTCA C	CCTGTAGATAAAACATG GTACCAGGTC	CCTTTCTTGAT GGTATCC	TCCTTTCTTG ATTGTATCC	0,0	omy24	245476 87
Omy_R AD5599 7-10	A	C	CATTTTCTACCTGCAGGCTGC	AGCCTACATACATAAAG CCAACA	AGGCTGCAAT GTTT	AGGCTGCCAT GTTT	1,5,0	omy24	274058 86
OMS00 052	T	G	TGCGTTTTTCATCCCAATCAT TCAC	GGCATCAGGCTCTTCTT CCT	CTTCCTTTTGA GAATAAT	CCTTTTGCGA ATAAT	0,0	omy24	281749 67
Omy_10 7336- 170	C	G	GCCCTCTCACTCATGACATC AAC	GCTCCAGCCACTCGCA	CACTCCTGGG TGCAGAA	ACTCCTGCGT GCAGAA	0,0	omy25	300174 55
Omy_R AD5975 8-41	T	C	GGCCCCCTTCTTTTCAGGAAT	CACACACTCAACGGGTC AGT	TGATTGCTACT GAC	TGATTGCCAC TGAC	0,0	omy25	402193 56
OMS00 174	A	C	TGACTAACTATGCAGCCTGA AAGG	GGGATACTCTTGTAATA AACTGTTGGTTAGTA	CAAGAACAGG [AC]TAAATGT	CAAGAACAG G[AC]GAAATG T	0,0	omy25	473253 93
OMS00 071	A	G	CCGGAGTGACCTCACATTTG G	GCATCGTACAGTTCACC TACCT	CTTGTTTGAGC TTTTTCT	TTGTTTGAGC CTTTTCT	0,0	omy25	482484 74
Omy_pp ie-232	C	T	CTGTTTTAGATTAGAATGTTT TTGGTCAGGT	CTGAACATAGGCTTTCA TTTCAGACAT	AAATAGCGGA GAAAAT	AAAATAGCAG AGAAAAT	0,0	omy25	549909 25

Omy_R AD1403 3-46	A	G	GCAGGAGATTTATTTGGCCC C	ACCCTTGTGATCACATA CTGTCT	ATAGAGGAAT AGAC	ATAGAGGGAT AGAC	0,0	omy25	600640 21
Omy_R AD1186 -59	A	G	CACAGCCTGGATGTGGTTCT	ACAAGTTCCGGGAGTTT CCT	CCAGGACATC CAGG	CCAGGACGTC CAGG	0,0	omy25	702523 14
OMS00 002	A	C	TTTGATTTGATTTGTATCTGC TTCTT	CCAACATGCCTCACACA AAA	TGTTTTGCAGC GCTC	TGTTTGGCAG CGCT	0,0	omy25	823663 95
Omy_R AD4159 4-34	A	G	TGCAGGGTTATAATGTGTCT TTGT	AAATCTCGGGCTGAGGA ACG	CAGAGATACG TTCC	CAGAGATGCG TTCC	0,0	omy26	560047 5
Omy_cd 59-206	C	T	CGATTGGCCCAGATGTTTCC AT	GCTCCGTTGCATAGGTG ACT	CAACAATCGA AGGTAAAT	CAACAATCAA AGGTAAAT	0,0	omy26	802832 2
Omy_R AD1303 4-67	A	C	GAGTGATTCCCAGCCCTCC	TCTCTCCGTTGGCCAGA AAC	ATAAATCACA A	CTAAATCACA A	0,0	omy26	125370 39
Omy_ad a10-71	C	T	TCTTTGAGCGACAAAGTCCT TGT	ACCCACACATGAACGCA AAAG	CTTCCTGCGTC CAA	CTTCCTGCAT CCAA	0,0	omy26	183154 20
Omy_as pAT- 123	T	C	GCCCATTTCCTGATGCTGT GA	AGGAGACCACTCCAAA GAGAACT	CCTTCCTAGGC AGTCAG	TTCCTGGGCA GTCAG	0,0	omy26	192895 82
OMS00 014	T	C	CTTACACACAAGGGCTTCAT TCTG	GATGTCTCTGGGTGGTT GTCA	TGATTTGATG AATTAACTT C	TTGATGAATT GAACTTC	0,0	omy27	629752 6
OMS00 015	A	T	TCAGACCCTATTTTGGCAC AAGT	GTCTAACTGATCCCACT TCTGCAT	CAAGTCACAC TTTAAATGAA	CAAGTCACAC TTATAATGAA	0,0	omy27	237309 45
Omy_hs p47-86	T	A	CACATTAAGCACTCCCAGGG A	TTGCAAAGGCCAAACAG CATT	CAGGAGTGTA AATGTTT	ACAGGAGTGT ATATGTTT	0,0	omy27	279444 94
OMS00 090	T	C	AGGGCACAACACCACTCTAA ATT	TCGAAAAGCAACATCTG TCTCAGT	ACAACCACAC AAGATT	AACCACGCAA GATT	0,0	omy28	424599 7
Omy28_ 1160795 4	G	A	TGACACTGATCACAATGGTG AAAT	TAAACTGGAAGGAGAG AGCAAAAT	TGTGGGCTGC GAACATACTC A	TGTGGGCTGC AAACATACTC A	0,0	omy28	116079 54
Omy_R AD5245 8-17	C	A	ACGTGTCCCTGAGGATGGTA	AGCTCTAGGTCTGGGTC CTG	ATGGCCCC[CT]AAGAACCC	ATGGCCCA[C T]AAGAACCC	0,0	omy28	116097 94

Omy_G REB1_0 5	T	G	TGGGCAGATATGGAAGAACG G	ACCTTCTAAATGGCCTC TGTGT	CGGTGGCTCT C	CGGTGGCTCG C	0,0	omy28	116180 27
Omy28_ 1162524 1	A	G	CAACATTTAGGGAGAGGTTG CTAT	ATCATCAAGTTTGCCTA CGACAC	CCTCCTCCCTA TGGTTGTCTC	CCTCCTCCCT GTGGTTGTCT C	0,0	omy28	116252 41
Omy28_ 1163259 1	G	A	GTAGAGGCCAAAGGCTTGAG	TGCTCTTATTACCTTCCA GACTCC	TGAGAAGAAC ACAGAGG	TGAGAAAAAC ACAGAGG	0,0	omy28	116325 91
Omy_G REB1_0 9	T	G	CCAGTGGCAACCTCAGGTAG	GACTCCAGTCACCCAAG TCA	TCAATGGAGA	TCAAGGGAGA	0,0	omy28	116416 23
Omy28_ 1165885 3	A	C	CAACATATGACCACTCGAAA ACTC	ATTAATCACACCGTGAG ACTCCTC	TGGTACAGAC ACGCACTAGC A	TGGTACAGAC CCGCACTAGC A	0,0	omy28	116588 53
Omy28_ 1166757 8	T	C	ACAGTAAACCCATTTCAGGCA TAGT	TTATCCTCTCAATCCAC ATCAAGA	GTATTGATCCT GTGGGAGACA	GTATTGATCC CGTGGGAGAC A	0,0	omy28	116675 78
Omy_R AD4708 0-54	A	G	TCAAAACCTGCAGGACTTGG A	TGGTTATATCTACAGTA CAGTTCGT	TGCAAGACTT AAAACGA	TGCAAGGCTT AAAACGA	0,0	omy28	116679 15
Omy_R AD1570 9-53	G	A	TGCAGGACTTGGATAACACA GA	TGGTTATATCTACAGTA CAGTTCGT	ATGCAAGGCT TAAA	ATGCAAGACT TAAA	0,0	omy28	116679 15
Omy28_ 1167111 6	C	T	AATTTCCCCAAATTTGAAAC TCTT	GTGTACATTGTCAGGCA GAAACAT	CTGGTGAGAA CAGGAATTAC C	CTGGTGAGAA TAGGAATTAC C	0,0	omy28	116711 16
Omy28_ 1167662 2	T	G	CGAATGCACTGTAGCTCATT CTAA	GCAGTAGAATGTCTCGC AAATACA	ACATGTCATTT ATTGTTATCT	ACATGTCATT GATTGTTATC T	0,0	omy28	116766 22
Omy28_ 1168320 4	G	T	CAAGAAAGAAACAGATGTTG TCCA	TTGTGACTCAAATCTGC AACCTAT	ATGTAAAAAA GGGCAGAAAA	ATGTAAAAAA TGGCAGAAAA	0,0	omy28	116832 04
Omy28_ 1177319 4	A	T	AGTTTGACACCCCTGTACTA GAGC	GTCTAACAAGCTCTGGG TGATTTA	GCAATTTTTTTA AAATTACCGC	GCAATTTTTTT TAAATTACCG C	0,0	omy28	117731 94
Omy_R AD7606 0-20	C	T	TGCAGGGTGTCTAGTATTGGG	TCCCATGCAAATTCCAA ATGCT	GGGCGCTGTA GGCAA	GGGTGCTGTA GGCAA	0,0	omy28	215613 03

OMS00 129	C	G	GGAGATGATGAAATAAAAAAT TGAGGAAAAGATGA	TGTCTGGTGAATTATCG CAAATAACCA	TTGAACAACA AGAAAAA	TTGAACAACA ACAAAAA	0,0	omy28	225311 25
Omy_R AD4645 2-51	A	G	TGCAGGTAAGACTTGATCTG GA	TGACTCCAACCTAAGTG CATGT	TGAAGTCAGA AGTT	TGAAGTCGGA AGTT	0,0	omy28	275166 10
Omy_97 660-230	C	G	TCAGTTATGTGTAATCTCATT ACCTCTCCAA	AACAGAAAAGGTCTCA ATGTATTTTTTGCA	ACGTAACCTTG TAGCGTTTT	ACGTAACCTTG TACCGTTTT	0,0	omy28	357429 46
Omy_im pa1-55	C	T	CGCTGAGAGGATTGTCAA	TTTTCTTTGTTTCAGTCTT CTGTCTCTG	CGAGATGATG CGTCTACA	CGAGATGATG CATCTACA	0,0	omy28	363905 09
Omy_B AC-B4- 324	G	T	CGTACTTTTCTTTTACAAAAT TAAGTGGAGGAT	GCCTAATATTGGCCTAA TGTCCTTCA	CATTGCCAAA TACG	TACATTGACA AATACG	0,0	omy29	145286 35
Omy_O myP9- 180	C	G	CTGGATGTGTAGTATCGGTG GAAAA	CACTGGGCACCTCTGAT CTC	CTGTAGTAGT CCCCATTGT	CTGTAGTAGT CCGCATTGT	0,0	omy29	156734 14
OMS00 164	T	G	CAGAGGAGAGGAGAGCAAA ATACTT	ACAACCTACTCATTGAA ACTCATTGGA	CCAGATTCAA TTAAATTTA	CAGATTCAAT TCAATTTA	0,0	omy29	172496 54
Omy_cr b-106	G	T	GCTCAAAAAGATTCTGCCAA ATTCACA	ATTACAATGAAAGTACT TGAGTGTTTATGCAAA	TTGCAATGCG TCTTT	TTGCAATGAG TCTTT	0,0	omy29	307013 74
Omy_11 8205- 116	A	G	CTGCGGTGGGCTACACA	CGCAGCTGCGGATGAG	CTACTGAGGC TGAGTGCT	TACTGAGGCC GAGTGCT	0,0	omy29	332671 27
Omy_sS OD-1	T	G	GCCGGACCCCACTTCAA	CAGACTAACCGAACAGC ATCAGTGG	CCACAACAAG ACCC	CCACAACCAG ACCC	0,0	omy29	398022 21
Omy_R AD1957 8-59	A	G	GGTTGGACACCTCCTGGTTA	TCAACCAAGCAACAGAT TATAGCT	GGTTAAGAGT ATTC	GGTTAAGGGT ATTC	0,1.8	NA	NA
Omy_R AD4052 0-48	T	G	TGTTCATCTGATCAGCTGTCA G	ACACGTCGGTCTTCTTC TCC	GTCAGATTGC GCTG	GTCAGATGGC GCTG	0,0	NA	NA
Omy_R AD7814 7-27	C	T	GCATTTTAGCCCTCCCAAAG TC	CCTTCTTCCAGTTGTAA AACCCA	CAAAGTCCCA GAGA	CAAAGTCTCA GAGA	0,2.6	NA	NA
Omy_R AD619- 59	T	C	CATGGAGAAACAGACCCGCT	TGCTGTGTGTGTATCTG GGG	TGCTGGATCC CCCA	TGCTGGACCC CCCA	0,0	NA	NA

1004	Omy_R AD9871 5-53	T	G	CGTAACGGGGAGCTGATCTG	GCTGGTAAAATGCTGAG GGG	CAGGACTTCT CCCC	CAGGACTGCT CCCC	0,1.9	NA	NA
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Assay	A1	A2	Forward	Reverse	A1-Probe	A2-Probe	Allele Corrections	Koop GCA_ 00287 2995.1 chromosome	Koop GCA_ 00287 2995.1 snpcordinate	Chi6 GCA_ 00283 1465.1 chromosome	Chi6 GCA_ 00283 1465.1 snpcordinate
Ots_SEXY3-1	X	Y	GGTCTTGCAGTCAGGA GAGG	CCAGGTGGTGAAG GTAGGAA	NA	TCAGCGAA GTGGAGAT	NA	NA	NA	NA	NA
Ots_myoD-364	T	G	GTGTGTGTGTGTGTGT GTCATCGT	TTTACACATATACA AAAATGGTCCTCTA TTGTCAT	TCATCTTTT GTTATTTCC TTG	ATCTTTTGT TCTTTCCTT G	0,7, 0	Ots01	10134 698	Ots01	72997 84
Ots_u07-25.325	T	C	AGACAATCATGGTGTT TTGAGTCTTTCT	GCCTAGGCTTGATG GAGTCA	CCGCTTGAA AGTTTGA	CGCTTGAA GGTTTGA	0,0	Ots01	12626 231	Ots01	91987 21
Ots_110689-218	T	G	GTATAAACTAGAGTCC AGTGTTATGTTAATGT CTT	CATGGCAGACAAC AGTAGAGAATATG A	CACCAATCA ATTAATTAT T	ACCAATCA ATTCATTAT T	0,0	Ots01	15276 095	NA	NA
Ots_129458-451	T	C	TGGGACCCACATAAAG CAACTG	GACATAAGACCCAT TTAGCCCCTTTT	CATCTGGCA ATGCCTT	CATCTGGC AGTGCCTT	0,0	Ots01	15863 134	Ots01	11080 640
Ots_crRAD21115-24	C	T	TGCAGGTGGGACTTAA ACACA	ACCTGTGGCAACGG TTGA	CACACACAT GCACG	CACACATA TGCACG	0,0	Ots01	27727 076	Ots01	23499 747
Ots_Prl2	A	G	CCTGGTCTGTTTGTGAT CAAGATG	GGTTAACTCAAATA GAACATACTCTGAC ACA	ATGTATTGT TCATTTAAT G	TGTATTGTT CGTTTAAT G	0,0	Ots01	46840 237	Ots01	40924 770
Ots_IkaroS-250	G	A	GAGGCTGACTTGGA TTGC	GGCCTGTCAGCCAA GGA	ACAGAAGA TTTTCGGCT GC	ACAGAAGA TTTTCGACT GC	0,0	Ots01	50283 457	Ots01	45385 098
Ots_107806-821	T	A	TGCAGTGCTGAATTAG AGATTAATTTTTGTG	CTCCCTTGCTTTTG GTCATTGG	CAAAGAAA ATCAAAATT T	CAAAGAAA ATCTAAAT TT	0,0	Ots01	56999 741	Ots01	51993 828
Ots_127760-569	C	T	CTGCTGGCGCAGACAT G	CGTTATAGAGGATA GTTTGGAGGAAGG A	CCGGTTTAC CGATTTG	CGGTTTAC CAATTTG	0,0	Ots01	70642 723	Ots01	65444 532
Ots_crRAD23631-48	G	A	GCCATATCCCGGGGCT TG	TGCCTCTGAGCACT GACTG	GGGCTTGGG GGCAT	GGGCTTAG GGGCAT	0,0	Ots01	73494 810	Ots01	68018 794

Ots1_728 58599	A	G	AACCATTGTTCTTGTAT TCCTGCT	GTAAAAAGACGAA TGAAAGGATGG	GGTGGAGG GAAAAAGC AGTG	GGTGGAGG GGAAAAAGC AGTG	0,0	Ots01	78671 220	Ots01	72858 599
Ots_unk3 513-49	C	T	TTTGAGTGAGTCACTG CACCAA	CAGCTCCACAGTGT CACCAT	AGTGCGAA GAACC	AGTGCAAA GAACC	0,0	Ots01	81595 603	NA	NA
Ots_crRA D74766- 28	G	A	GCTGACCACCGACCAC AG	AGCTCTGCAGTAAC AATGGGA	AGACTGGTA AAAG[AT]	AGACTGAT AAAAG[AT]	0,0	Ots01	88530 200	NA	NA
Ots_crRA D57376- 68	T	C	TGCAGGCATCATGCTT AATAACT	ACGTGACACAGGTC TGGG	ATAAAGTGT GTTAT	ATAAAGCG TGTTAT	0,0	Ots02	99375 16	Ots02	54571 149
Ots2_424 05643	G	T	GAGAGAGTGCATTCTT CATCAAGTT	TCAGTACAGGGTTT TTCCCAAT	CAGGTTGTT GGTTGTT	CAGGTTGT TGTTTGTT	0,0	Ots02	19918 478	Ots02	42405 643
Ots_crRA D75581- 70	A	G	ACACATGGCTCGTCTG CA	GGAGCTCAGGGTGC AGGA	GAACTTAAA ACACT	GAACTTGA AACACT	0,0	Ots02	20994 028	Ots03	27546 510
Ots_crRA D46081- 56	C	T	GCAGGGTCTGTGTGGG TT	ATGAGGACACTCCG CCCA	GCACCACTG GACCC	GCACCATT GGACCC	0,0	Ots02	24120 152	Ots02	34735 614
Ots_nelfd -163	A	G	CTCACTGCAAATCCAA CTTCATCAT	CCACTACATCCTCA TCCAAGGTT	ACCCACCAG TGTCATT	CCACCAGC GTCATT	0,0	Ots02	27801 414	Ots02	31069 128
Ots_crRA D27515- 69	T	A	CAGATGGTGCAGGCCG AA	ACTCGTTGTGATTC CAGCCA	GCATTTTAA AAATC	GCATTTAA AAAATC	0,0	Ots02	31417 111	Ots02	27211 812
Ots_1283 02-57	C	T	GGTTGCAGGGCAGAAC TGT	ACCCATCCAATAAC CCATTTTCCTT	CCTGCAATA CGACCAAC	CTGCAATA CAACCAAC	0,0	Ots02	34870 376	Ots02	23632 129
Ots_crRA D78968- 46	C	T	CCTGCTCTGTGTCTGG GC	GTGAAGACGACCCC GGTG	AG[CA]AATC [CA]CACAGC	AG[CA]AAT T[CA]CACA GC	0,0	Ots02	35362 682	Ots02	23276 932
Ots_crRA D11620- 55	C	T	TGGGATAGAACAGGA GCTTAAACA	TGTCTTGGTCGCGC AGTT	AGAAGCCC AGCTCC	AGAAGCTC AGCTCC	0,0	Ots02	36061 414	Ots07	43120 003
Ots_1054 07-117	T	A	TGTGTACATCCGCGTA AATATTGAAGATAA	CTGTGAGCTGCTGC AAACC	CAGGTTAGG AATGGTTG	CAGGTTAG GATTGGTT G	0,0. 2	Ots02	37138 862	NA	NA
Ots_LWS op-638	T	C	CAATTACTCTTTCTCAG CCCTGTGT	GCGGTAAGATGCA GTTTTACATGGA	TTTAACAAG AAAATTATA CATTTTC	CAAGAAAAG TTATACATT TC	0,0	Ots02	44036 657	Ots02	13620 416

Ots_crRA D69327- 53	G	T	GCCATTTGACCAACGG AGC	ACTCATACAGTATT TCCGCCTGT	ATAGGAGA ATTGGA	ATAGGATA ATTGGA	0,0	Ots02	52051 529	Ots02	47082 99
Ots_stk6- 516	C	A	TGTGTTTtaggATTGAA CTGACCATGTT	GTAAACTCCACCTG CAAGAAGGA	AACATAAC GGACTCCC	TAGAACAT AACTGACT CCC	0,0	Ots02	53397 399	Ots02	38145 27
Ots_cgo2 4-22	T	C	AGGTCCTCTGTCGCAC CTA	GGAGGCGAGGTCT GGTG	CCAGATGA ACAACCTCA C	CCAGATGA GCAACTTC AC	0,0	Ots03	95886 38	Ots03	12326 537
Ots_NOD 1	C	G	GTGCTGCAGGAACCAT GTG	CTGTGTGGACTGCT GTCTAAGG	CCAACGGC GACTTG	CCAACGCC GACTTG	0,0	Ots03	95985 76	Ots12	23568 768
Ots_1134 57-40R	C	T	CCCAAGTGGTGAGTGT CAGT	ACTACAACAGGTGT TGATAATAGAATCA TTCTC	CCCT[AG]TT CTCCAATCC ATAT	CCCT[AG]TT CTCTAATCC ATATG	0,0	Ots03	17561 381	Ots23	94474 90
Ots_TNF	C	T	CCAAATCCTCATCCCA CACACT	CCGTTGCACTTGAC CCTAAAC	CTGGCTGTA AACGAAGA	TGGCTGTA AACAAAGA	2,0	Ots03	20344 952	Ots03	21402 560
Ots_1123 01-43	T	C	GCATGGCTGCCCTAGA ACA	TCAGAACATTTTCCT TCAGCTTCGT	CGTCGCATT CAGC	CGTCGCGT TCAGC	0,0	Ots03	33896 915	Ots03	284X XXXX
Ots_u07- 20.332	A	C	CGCGAGTTAGCTCGAA TATTATGATTTT	TCAAGCTAGCATAG CAACTTCATCAA	ACCATTTGA TATAACTGC GTTAG	CATTTGAT ATAACGGC GTTAG	0,0	Ots03	39126 796	Ots03	33660 993
Ots3_348 94254	T	C	TGATATATTTTGCTGC AATGATCTG	AGAGGGAAGGTGG ACAATGTT	TAAC TTACA GTC	TAAC TTAC AGCC	0,0	Ots03	40436 224	Ots03	34894 254
Ots_1189 38-325	C	T	ATTTTCAAACAGGCAT TTATCATTGGTGAA	GGTCTGTCCCTCAT TCTTTGCA	AGAGATGC AAAGTGGA GTT	AGAGATGC AAAATGGA GTT	0,0	Ots03	48430 243	Ots03	43610 834
Ots3_570 55518	T	C	TTAGCAGGCGATCTAA TTCTGATT	ACACGCTAACTGCT GTATTCTCTG	TGATCATAT CTCGTTCAG T	TGATCATA CCTCGTTCA GT	0,0	Ots03	61138 782	Ots03	57055 518
Ots_1008 84-287	T	C	CGGAAGACCAGATTCT CCAAGAGTA	CGACCAAGTAGCG GCACTT	ATAGAACTA CAATTCACA TATAT	AACTACAA TTCGCATAT AT	0,0	Ots03	72097 697	Ots09	34802 127
Ots_9485 7-232R	T	C	GGCACTCTCCCTGGCT AGA	CCCCATCACTTCTC TGGCTTTAAAT	CAGGATAAT AACAAACA AG	CAGGATAA TAACGAAC AAG	0,0	Ots03	2XXX XXXX	Ots03	27503 532
Ots_1072 85-93	T	A	GCCCTTGTGACAATGC ACTGTTATA	AACATACACCAATA CTTAGGTCTAGACA GT	AAGTAACGT ATCAAATGG C	AAAGTAAC GTATCATA TGGC	0,0	Ots04	66909 09	Ots04	58529 87

Ots_Ots3 11-101x	A	-	AAATGAGGCCGTCCTT TACACT	GCAATACAAGCCCT TGATAATGAAGT	CTGAGATCA CTTTGAGCA C	ACTGAGAT CACTGAGC AC	0,0	Ots04	20118 827	Ots04	40706 52
Ots_1031 22-180	T	C	CAAACGCGCACTCACA CACA	TCACAATGGTACGA TTTTACGACTCAA	CATCAACAC AATCTGC	CATCAACA CGATCTGC	0,0	Ots04	22443 878	Ots04	16781 457
Ots4_409 42276	G	A	ATTAGTGCATATGAAT CGGGCTAT	CCAAAAGGTTGAAC GTACATATTG	GGAGTCAG ATAC	GAAGTCAG ATAC	0,0	Ots04	48716 473	Ots04	40942 276
Ots4_416 38710	G	A	CAGCAGCTGTTTATGA CTGACTTC	CTCGTGTAACCGAT GTGAAATG	CCTGAGATT AGG	CCTGAGAT TAAG	0,0	Ots04	49353 548	Ots04	41638 710
Ots4_423 78741	C	T	CAGTTTAAGTGTTACC ACCACGAG	GTGCAGGTGAGCGT TAACG	AGATGAAC ACCAACTGG CCGG	AGATGAAC ACTAACTG GCCGG	0,0	Ots04	50072 182	Ots04	42378 741
Ots_9650 0-180	G	T	CAGGTCTGGTCTACAT CGAACAC	GATCATGTCAGATA GGATGCTGAAAGT	AAAACAAA TCATTTTTC G	AAAAACAA ATAATTTT CG	0,0	Ots04	53468 710	Ots04	45636 897
Ots4_649 78818	C	A	AGAACCCATGCTTTCA GTACACTT	AAAATGGACAGAA ATGTATCGCTC	TCAAGTGTT TCCTTTATT TTG	TCAAGTGT TTACTTTAT TTTG	0,0	Ots04	71212 183	Ots04	64978 818
Ots11_11 925999	G	T	TTTATATTCAGACATTC GCCAAAA	GATTGTCACAGTAA CATAGAAATGGTTT	CATTTAAAA TGGTAAAA ATCA	CATTTAAA ATTGTAAA AATCA	0,0	Ots04	71231 942	Ots11	11925 999
Ots_1054 01-325	G	T	GAAGTGAAGCGGCTGCT G	CGCCTCCTGGTGTC TATCCT	CCCGGACA AGATGAGA CAG	CCCGGACA AGATGAGA CCG	0,0	Ots04	1XXX XXXX	Ots04	12543 170
Ots_HSP 90B-100	C	T	CACCTTAGTTCCACGC AACATG	CTGCGTGTATTGTA GTGGTGACA	TCTATGGTG TGATTCAAT	TTCTATGGT GTAATTCA TT	0,0	Ots05	76415 77	Ots05	68666 92
Ots_1116 81-657	G	T	CTGAGCTTTTTCAACTT ACTTGTTGGA	GGCGCAGCAGCAA CTG	TAGCGCAA ACCCCGAAC C	CGCAAACA CCGAACC	0,0	Ots05	94397 42	Ots05	85399 47
Ots_1286 93-461	C	T	TCAATGTTTCATCAATG CACTTCCTGTA	GCCTGCAGGAGAA GGTAGAGTTA	CTGGTACCC A	CTGATACC CA	0,0	Ots05	15919 055	Ots11	29099 828
Ots_crRA D66330- 60	G	T	ACTCTCCAGAAAGGAT TCAGAGA	TCCCAAAGCATCCT GCCA	AGAGAGGG GTCAAA	AGAGAGTG GTCAAA	0,0	Ots05	17933 258	Ots05	14208 918
Ots_u211 -85	C	T	TGGTGAGAGCAGCTTT AAATGTCTT	ACCCATTCTTCTGT CTGGTTTAAGC	TCCCAAAGT CGAGTGTG	CCCAAAGT CAAGTGTG	0,0	Ots05	29079 237	Ots05	34253 781

Ots_1209 50-417	A	T	CAGACAGGTCACCATC ACACT	TGGTGAAGCTGTAG GAGAAGGA	CTGGACCAG AACTCTGA	CTGGACCA GATCTCTG A	0,0	Ots05	31561 077	Ots05	18867 357
Ots_HM GB1-73	G	T	TGCTTCAGTGAAAATA AGCGTGAGA	GTCGAGCGGTATGA ATACTTTCTGA	ACTGTATAT GTTACGTTT TC	ACTGTATA TGTTAAGTT TTC	0,0	Ots05	39590 504	Ots05	27531 907
Ots_BMP 2-SNP1	C	T	ACTGCCACAGACACGA ACTC	GCCACTATCCACTC GTTCCA	CCCACCTCG CTGAAGT	CCCACCTC ACTGAAGT	0,0	Ots05	42344 711	Ots05	30442 475
Ots5_447 95073	C	T	GCACTGTATACAAAAT CGTGTGGT	CATTAAGACAGACA ATGCCCAATA	TTTTTGTGT CCGCCATGA ATT	TTTTTGTGT CTGCCATG AATT	0,0	Ots05	54248 412	Ots05	44795 073
Ots_Cirp A	C	T	GCTGTGATTGTGCTCT AAAGACATG	CTCCCACCTTAGCAT TCCTACCTT	CAGTTCTGT AATGCATT	CAGTTTTGT AATGCATT	0,0	Ots05	56708 050	Ots05	47621 841
Ots_1272 36-62	T	A	TGGAGAACTTGCCTG AATGTGAAA	GCTGTTGGACCTTG ACTTTAACAAATT	TCTCTTATC TGAGTTCTG C	CTCTTATCT GTGTTCTGC	0,0	Ots05	61558 078	Ots05	52450 716
Ots_u4- 92	T	C	ATCCAAGGAGCCCCAT TAAAGATTT	CGTACCAGAGTTGT AGAAGCATCT	CTGTGTTGA ATTTAACAT AAT	TCTGTGTTG AATTTAAC GTAAT	0,0	Ots05	62235 849	Ots05	49324 631
Ots_unk9 480-51	G	C	CAAATCAGAACAAAAC CTCCCACAA	GGAAGTCTGTCTGA ATGGTTGTCTT	CTCCCACAA ACCC	TCCCAGAA ACCC	0,0	Ots05	68359 607	Ots05	58885 154
Ots_1095 25-816	C	T	GCCAGATAGTAGCGTA CATCATGAG	CTCCCCATGTCCCT GAGTCT	CATGAGGC GTTCCGC	ATGAGGCA TTCGGC	0,0	Ots05	86382 977	Ots05	70296 658
Ots_crRA D47297- 55	T	C	CTCCCTGTTCGCTAGC CG	GGACGACCAAAGG TAGAACCC	TAGCCGTCA CCGAT	TAGCCGCC ACCGAT	0,0	Ots05	91281 495	Ots12	48785 08
Ots_SClk F2R2-135	A	T	CCAAATACAGACCAGC TACTTGTGT	CTTCAAGTCCCTGA ATAATGGTACGT	ATTCAAAGT CAAATTTT	ATTCAAAG TCTAATTTT	0,0	Ots05	XXXX XXXX	Ots05	60109 411
Ots_crRA D13725- 51	C	A	TGCAGGAGGAGGAAG GCA	AGAGCTGCCAGGTG GAGT	GAGGCCCC AGATTC	GAGGCCAC AGATTC	0,0	Ots06	12958 480	Ots06	10662 607
Ots6_109 04949	C	T	AAATCACCCCATTCT TTTGTG	GTAGAAAGGTGAT GTGCATAAGCA	CCTTTGTCA CCGCTCATC AGC	CCTTTGTCA CTGCTCATC AGC	0,0	Ots06	13203 375	Ots06	10904 949
Ots6_335 05144	T	A	CCCACCATACAATAAA GGCATGT	ATCTCTCCCATAAA CAAATACCCA	AACATATGA GTTGTAATG CCC	AACATATG AGATGTAA TGCCC	0,0	Ots06	14379 077	Ots06	33505 144
Ots_OTS MTA- SNP1	C	T	GCCGAAAAATAAGCG ATTAGTGATGA	GCCCCATGGTAAAC CTAATTAACCT	AATTGCCTC ATTGGGTG	AATTGCCT CATTAGGT G	0,0	Ots06	21997 348	Ots06	20396 609

Ots_Met A	T	A	GATCATTATCAAGAC TATAGGCTATGGATAC G	AGTTGAGTTAAGTA ATTGGTAATTAGCC TGTT	CCTTAAGCA TATTTCT	CCTTAAGC GTATTTCT	1,6, 0	Ots06	21997 591	Ots06	20396 849
Ots_ZR- 575	G	A	GCCTACCAGAAAGTAC CAATTGTGA	ACTTTTCACTGTCC TATTACAATTAGTA TTTGTGATAT	CC[GA]ACAC AATTTTGT	CC[GA]ACA TAATTTTGT	0,0	Ots06	29177 962	Ots06	27576 884
Ots_crRA D46751- 42	C	T	CAGGAACCTGCTTTAA TGCTCT	GCTTCTGCAGGGGG ACAA	TTTCTACTT AGTAA	TTTCTATTT AGTAA	0,0	Ots06	35130 855	NA	NA
Ots_9490 3-99R	G	T	CCGTCTGAGTAGGAGG ATCAATACA	TTTGGATCCAGCTC TCCGTATAGA	CAAACCAG CAAACAT	ACAAACCA GAAAACAT	0,0	Ots06	36076 090	Ots06	35539 172
Ots_1076 07-315	A	C	GTGATGAGAGGTTTCC GGAAAATCT	GTGTTCTGGATTCC ATTGTGCAAA	ATGGGAGA CAGATAACT	ATGGGAGA CATATAAC T	0,0	Ots06	36480 917	Ots06	36761 804
Ots_FGF 6B_1	A	C	GAGACAAAGGTTTGCA GGTTCATG	GGGAGCCATGCACT AATATATTGGA	CCTGTTATC AGACCCAA AT	CTGTTATCA GCCCCAAA T	0,0	Ots06	37585 917	Ots06	37250 687
Ots_u6- 75	C	T	GAAAAAGTAAAGTAA AAGTAAAGTATTATAC CACTAAAGACAAT	GATCCACACTGTTG GTCTACTACAA	TTAGTCAAC TGTTGTTTT T	TTAGTCAA CTGTTATTT TT	0,0	Ots06	50534 743	NA	NA
Ots_crRA D73823- 60	T	A	GCAGGAAGCAAAGTTC GGTG	AGCAACTCATCGCG TGGT	GCACGATG[CT]AGAAC	GCACGAAG [CT]AGAAC	0,0	Ots06	52896 472	Ots06	53325 628
Ots14_54 53033	G	A	ATTATTCAAACAGAGA TGGCGAAA	GAGGAGGTTTGCAT AGAACATGAT	TCTCTAAAA AGGTACAGT ATA	TCTCTAAA AAAGTACA GTATA	0,0	Ots06	55710 320	Ots14	54530 33
Ots_u07- 53.133	C	T	AGCTAGGCTGTAAATG CAAGGAT	CAGTGCTTTCAATT CATGCTGTCAA	TAACACATG TTGGAGGTC	AACACATG TTAGAGGT C	0,0	Ots06	58232 274	NA	NA
Ots_GDH -81x	C	-	CTTTTCTGAATTAGTGC TGTGCTTGT	CCAACCTTCTTCAAC TCTGTCAGTGA	TGTTACGGG ACATACT	TCTGTTACG GACATACT	0,0	Ots06	58688 462	Ots06	585X XXXX
Ots_GnR H-271	C	T	CAGATGAAAAATAAAT AATTGGGCCATTAGGA A	CAGAGAGACTGAG ACCATATGATGTAG T	CAATGAATA CAATATCTA ACCTAAT	AATGAATA CAATATCT AATCTAAT	0,0	Ots06	65879 558	Ots06	65660 176
Ots_1159 87-325	T	G	GGAGGTGTAGTGAAAT GGGAAGAT	GCATTCAGTGAACC AGTAGTGCTAT	ATGCATAAA AGGTAATTG TG	ATGCATAA AAGGTCAT TGTG	0,0	Ots06	66387 317	Ots06	66193 572
Ots_1053 85-421	A	G	GACTGTCTTGGAACCG TTGCTA	TCCCGGAACACACC AATGTC	CCTCCTGGG TATATCG	CTCCTGGG CATATCG	0,0	Ots06	73075 541	Ots06	73012 849

Ots_map Kpr-151	A	T	TGTTGTCTCGGACTGC ATGAC	GAAGGCACAGAGA TGAAGGACAT	CGTATGTGC AATGCATG	CGTATGTG CATTGCAT G	0,0	Ots07	35931 75	Ots07	54601 43
Ots_map K-3'-309	T	G	GGCCACTGTCATAGAA TTAGGCATT	CGTGACCCTTGTA CTGAAAAGC	ATGCTATTA AATGAATAT TC	ATGCTATT AAATGACT ATTC	0,0	Ots07	36086 28	Ots07	54800 42
Ots_1287 57-61R	A	-	CGTGTCCGGCTTCTTTT ATTTTCATT	GATGGGTATGTTAA TCATATTACCAGCG TAA	TTGTGCATT TTCCCC	TGTGCATTT CCCCC	0,0	Ots07	66438 30	NA	NA
Ots_1318 02-393	C	T	TGATTGTCTCATGGCC AATTGTCA	TGTAAATTCCACTT GGCAATCTTTGG	TGTTTCGAGA ATGAAGAT GAGTAA	TCGAGAAT GAAGGTGA GTAA	0,0	Ots07	35733 706	Ots07	25665 994
Ots_1314 60-584	T	C	CCTATTTTTGATAGGTC ATAGTGAATGGGATAG	CTGTACTCCTCCAT TCCTTTTCACT	CTATCAAAG CAATACATT G	CTATCAAA GCAGTACA TTG	0,0	Ots07	42945 148	Ots07	31130 645
Ots_CHI 06048618 _5222	T	G	GCAATTACCCATGACT CTGTGA	GCCAAAAAGAGAC CGAATCA	ATTGTGCTT ATCACA	ATTGTGCTT AGCACA	0,0	Ots07	50130 039	Ots07	38043 969
Ots7_509 97124	G	T	AGATCAAGCTTGCTGA CTTCG	CATACACCACACTG TATTTTGCTG	GGGCCTTCG GGGTGCCTG TCC	GGGCCTTC GGTGTGCC TGTC	0,0	Ots07	62901 252	Ots07	50997 124
Ots_nram p-321	G	A	GGCCATCTTTCAGGAC GTACAG	GCATGCTCTGCAAT ACGTTGAG	AAC[GT]GGC ATGAACGA CTT	AAC[GT]GG CATGAATG ACTT	0,0	Ots07	63847 460	Ots07	51336 125
Ots7_514 09415	T	C	ACAACTAGTCATCGTG GAATCTGA	ACATGCTAAAAGA AAGGAATGAGG	TGGTCTACT TTGTGC	TGGTCTACT TCGTGC	0,0	Ots07	63921 389	Ots07	51409 415
Ots9_289 75221	A	T	GCCTGCCCTACTTATCT CTTATCA	ACCTCTTCACTGTA AAATTTGCTG	TTTGCCAAA GAGTTCAGA TAC	TTTGCCAA AGTGTTC GATAC	0,0	Ots07	64709 638	Ots09	28975 221
Ots_arp- 436	A	T	GCCCTGGAGAAGTACG TTTTAACTAA	GCAACCATGTCAAC ATTGCACATAA	CTAGGTGAA ACTTTTTTT AAA	CTAGGTGA AACTTTTTA AAAA	0,0	Ots07	65259 187	Ots07	52672 820
Ots7_532 91035	G	A	TCAAATTGAATGTAGA CAGATGGAA	AAATAGCTCGCAAA GCTAACATTA	GCTAGCAA ACGTCGCCA	GCTAGCAA ACATCGCC A	0,0	Ots07	66039 124	Ots07	53291 035
Ots7_536 31522	A	G	CTTATCTCAAAGGAAT GGGAATGA	CTACAGTATTTGGA TGCAGCTTTG	TGAGTTTTT AAGGGGTT	TGAGTTTTT AGGGGGTT	0,0	Ots07	66362 189	Ots07	53631 522
Ots7_542 12944	T	A	AAACCACGGTATCCTT TATTCATC	TTCCTAACCCCAA ATATGTCTAAA	AATATATTT TTTATAGGC	AATATATA TTTTATAGG C	0,0	Ots07	66905 811	Ots07	54212 944

Ots_9707 7-179R	G	T	CCTGAACAAATACTTA ACGCTCCAGTT	GTAATAATACTTCA CACCATTGCCACTT C	TCACAAATG TATCCTAAA GC	CACAAATG TATACTAA AGC	0,0	Ots07	70085 972	Ots07	5632X XXX
Ots_1051 05-613	C	G	AGTACAAGTGCAGAGA ATGACATCATG	GGTGTTTTATTTTCC CATATATCTTTTAA CTTTAAGCT	CCGAGCTTG AGTTAGGA	CCGAGCTT GACTTAGG A	0,0	Ots07	77202 120	Ots07	63322 755
Ots_slc7a 2-71	G	T	CCATTCCCATCGGCAT CGT	GCAGCAGACACAC CGAAGTA	GTCTCTGAC GGTGTGCTT TC	GTCTCTGA CTGTGTGCT TTC	0,0	Ots08	86683 69	Ots08	61099 332
Ots_crRA D34397- 33	C	G	TGCCTAAACACTCCCA AGGT	GTTCCGTTTTTGTTC CGCGA	AA[GA]GTG CCTTCCCC	AA[GA]GTG GCTTCCCC	0,4, 0	Ots08	10116 544	Ots08	60161 043
Ots_u07- 64.221	G	C	GAGGATGACACTGTCC GTTTGT	CACAGTCCTTCGTA TTCACCTTGAT	ATCGACCCT GTCATTAG	CGACCCTG TGATTAG	0,0	Ots08	12242 337	Ots08	57998 143
Ots_1030 41-52	G	A	ACCACCCACCTCCTCA GA	AGACAGAGAAAGT CGGGACACT	CATCCTGCT GGACCC	CATCCTGTT GGACCC	0,0	Ots08	15518 130	Ots08	55669 454
Ots_OTA LDBINT 1-SNP1	T	C	CGCTGGGCATGGATGA GT	GGCCAACACTGCTA CTTCCT	CTACTGTTG TATTTTCTC	CTGTTGTGT TTTCTC	0,0	Ots08	16275 426	NA	NA
Ots_1017 04-143	T	G	ACTTCTTGAGCCAATC GGATGATG	CCAGAGATAAACTA GTGGAGGAGATCA	CTTAGACGT CAGAGGTC	CTTAGACG TCCGAGGT C	0,0	Ots08	21039 383	Ots08	51040 180
Ots_1022 13-210	A	G	CATTCCATGACAATGA TTGAAATCTAAAAACA C	GAGTATCTCAATTG CAACACTATGGTAT GT	CTGTATACA GTAAGAGT ATTAAT	ACAGTAAG AGCATTA T	0,0	Ots08	26477 291	Ots08	45XX XXXX
Ots_1291 70-683	C	A	AACCCTATGGGAACTC GTAGAACT	GCTAGGAGTTCTCA AAAGGGTTCT	ATTAGA CGTAGAACT AT	ATATTAGA ACTCGTAT AACTAT	0,6, 0	Ots08	47390 818	Ots17	11485 501
Ots_ppie- 245	C	A	TGTTTTTGGTCATGTAT TTTCTCTGCTATTTTT	GGACTGGAGCTGCT GAACATA	ATGTCTGAA ATGAAAGC C	AATGTCTG AAATTAAA GCC	0,0	Ots08	50220 193	Ots08	23260 130
Ots_Asn RS-60	T	C	CCGACGCCTCACTGAG T	TGGTTTTTCAGGTC ATGTTTTCCA	TGAGTCCCT GACCAGC	AGTCCCCG ACCAGC	0,0	Ots09	28000 47	Ots20	41784 960
Ots_u100 7-124	A	G	CGAAATAAGGGCCTGG TGTTTAAAA	TGTACCAGGTGGAA GCTTTGG	TGTCCTGTC CTCAGATCA	TCCTGTCCC CAGATCA	0,0	Ots09	11679 141	Ots09	94303 81
Ots9_161 15048	G	A	ATAGAGCTTTTGGTGT TTCATTCC	AGTGTGTGTACTGT GTACTGGCCT	CCAGTGAG ATGCTGTGT TGCA	CCAGTGAG ATACTGTG TTGCA	0,0	Ots09	19855 559	Ots09	16115 048

Ots_P450-288	A	G	ATGTCAATATATTTCA CTATAATGATTGGAAG CCA	CACTGAACTCGAAG CTGTTAGGA	CTATAAAGT TGGACAGTT GG	AAAGTTGG GCAGTTGG	0,0	Ots09	23147 965	Ots09	19138 835
Ots_102457-132	A	G	CCAGCAGAGACTGGGT TCAC	TTCCCTACCGGCGA AACC	TGGGGCAA CGCACAATT GGCT	TGGGGCGA CGCACAAT TGGCT	0,0	Ots09	26992 696	Ots09	23XX XXXX
Ots_110201-363	A	T	TTTTGGCTATTGAAAT TATACATTAAAACATG TAGCT	CCATGGCATCCTGT AAAGAACAACA	TTTTAAAA+ CTGGCATCC A	TTTTTAAAC TGGCATCC A	0,0	Ots09	29017 362	Ots09	25990 000
Ots_hsp27b-150	G	A	TAGGAGTTGGAAAGAC TGCACA	CCCATTGGTTCTTT GGTGTT	[CT]GATCTG GACCAGGCT	[CT]GATTTG GACCAGGC T	0,0	Ots09	67075 478	Ots09	52471 033
Ots_u1008-108	T	A	GGATGACTCCTACTAA TAGACGGATGT	AGGACAGGAAAGA AGCAGCAAATA	TTGGTAAAC CTGTTTATT GGTA	TGGTAAAC CTGTTTTTT GGTA	0,0	Ots09	75362 394	Ots09	62354 402
Ots_FARSLA-220	G	A	GTTCGTGGGATTGTTC AATGTTTCAT	CTTGGACAGGCTCA CATTACCATA	CCTTGGATG GGA	CCTTGGAT AGGA	0,0	Ots09	82809 166	Ots09	68274 964
Ots_hsc71-3'-488	C	T	TGCATCCATTTCATACC TGACCAATT	TTTGGTTAGGCACA CGATAATTTGC	TTTCCAATG GTATAGATA TGA	TTTCCAATG ATATAGAT ATGA	0,0	Ots09	XXXX XXXX	Ots09	37358 812
Ots_Thio	T	C	TTTTAAAAATGGAGAT AAACTCCTGACCTGAA	AATACCAAACCATG CCACTAATACCT	CAGTGTATT AGTCATTCT TA	CAGTGTAT TAGTCGTTC TTA	0,0	Ots10	71448 61	Ots10	56253 22
Ots_112820-284	C	T	CATAGATGTTTATATG AAAAACCTCCCACTGT	GCATCCAAAAAGA CGTGTGTGTTT	ACTCACACT CGAGTGACT	ACTCACAC TCAAGTGA CT	0,0	Ots10	18438 064	Ots10	16907 470
Ots_102414-395	A	G	GCCTACTGATAAATGT ATGACAGTAATGGA	CAATAACAAACAA GCTAGGAACAAAA GTGT	CACATAGTG TAGCTTTAC TAC	CACATAGT GTAGCTCT ACTAC	0,0	Ots10	21873 207	Ots10	20097 469
Ots_108007-208	A	T	CAGGCTTGTGTAAAGT AGGGAGAAA	CATTGGACAAGACC GGGTAGTC	CAGTTTCAC TTAATTTTA AAATG	TTTCACTTA ATTTAAAA ATG	0,0	Ots10	25247 839	Ots20	49711 42
Ots_129144-472	C	A	CTGTTAGTGCAGAAGA CGTAGCT	GCAGAGCTATTGAG CCAAGTTACAA	TGGGTCTCG AGCCTGTA	TGGGTCTC GATCCTGT A	0,0	Ots10	30541 697	Ots10	30704 018
Ots_crRAD10447-25	C	T	CCGTTGCAGGACTCAT CAGT	GCGTGGTTCAACAG CAGTG	AGCTAGCGC TCCTC	AGCTAGTG CTCCTC	0,0	Ots10	36392 206	Ots10	36699 634

Ots_crRA D18937- 60	G	A	GGCACAGCGACAGGA GTT	TGAGCTGGTGCGTC TGAG	CTCCTCAGG TGGGC	CTCCTCAA GTGGGC	0,0	Ots10	50635 181	Ots10	51017 984
Ots_nkef- 192	C	T	CATTTAGCAGACACTC TTATCTTAGTGTCA	CGAATGTCCACCTC AGATGTTACAA	AATAGGCC GACATCAA	AAATAGGC CAACATCA A	0,0	Ots10	51361 376	Ots10	51394 567
Ots_crRA D26165- 69	C	T	GGGCCACGGGGTTGTA AA	TCCCAGGATGCAAT GGGA	CTCT[GA]CC CCTGGAC	CTCT[GA]CT CCTGGAC	0,0	Ots10	52149 501	Ots16	20109 332
Ots_u07- 57.120	A	T	GGTTTGAGCCAATCAG TTGTGTT	CGGTCTAATGTCCA TTGCTCATGTT	GTGACAAG GTAGGGGT G	GTGACATG GTAGGGGT TG	0,0	Ots11	12974 126	Ots11	16346 192
Ots_1063 13-729	A	G	TTGTTCAATGGGCATT AATGCATGTT	TGCTTATGTGCAGA TACTTGAGACAAA	AAGAGTCC AGCGTTACT T	AAGAGTCC AGTGTTAC TT	0,0	Ots11	16517 829	NA	NA
Ots_1087 35-302	C	T	CCTTTTTCTTATTAGTT TTACTTCCCCAGAGA	CAATTCCATTCTTG ATTCTGTTTAACGG T	AAACAAAC AACGCCTCA TG	AACAAACA ACACCTCA TG	0,0	Ots11	17583 626	Ots11	21337 974
Ots_crRA D76512- 28	A	T	GCAGGGACAGGGCCCT	TGGTGCTGGGTGCT GTAC	TAAAAAAA TATAAA	TAAAAATA TATAAA	0,3, 0	Ots11	25425 413	Ots11	32328 258
Ots11_32 418659	A	T	CAATTGTAGCCCTCTA ACTTTTCC	ATACGACACACAA AGCCAATTGTA	AGCCAATTG TAGCCTTAG TGC	AGCCAATT GTTGCCTTA GTGC	0,0	Ots11	25519 430	Ots11	32418 659
Ots11_32 468959	G	C	AACACAGATCAAATGT TTTCACAC	AACACAGGAAAAA CAGAATGTTGA	GTGATAGTT TGATAGTTT TAT	GTGATAGT TTCATAGTT TTAT	0,0	Ots11	25588 829	NA	NA
Ots_crRA D9615-69	T	C	GAATGCAGGGCCAGG GAG	ACTCCCAGACCATC CAGCT	TATTGGTCA GGGAA	TATTGGCC AGGGAA	0,0	Ots11	25656 094	NA	NA
Ots_crRA D36072- 29	T	C	TGCAGGACCAACTTTC TCAT	GGCTGACTGGTGAA GGGG	AACCTGTGT GATTT	AACCTGCG TGATTT	0,0	Ots11	29147 224	Ots11	36019 633
Ots_crRA D61523- 71	A	G	GCCAAGTGATCAAGTG CTTGT	CCAGCAGTTCAGTT GCGG	CAGAGCAT GTGCTG	CAGAGCGT GTGCTG	0,2, 0	Ots11	35352 564	Ots11	41149 096
Ots_GPH -318	C	T	GGTGATAACAGGTGTT GCACCAA	TCAGGTGGTGGTGG ACAAC	ATCAAGCTG ACGAACCA	CAAGCTGA CAAACCA	0,0	Ots11	40844 253	Ots11	46176 184
Ots_crRA D20376- 66	G	A	GGGAGGCAGGCAAAA GGT	GGTTCACCACCAGC CTTCT	GGGA[TA]G GAGTATTT	GGGA[TA]G AAGTATTT	0,0	Ots11	41283 424	Ots11	46457 753

Ots_zn59 3-346	A	T	CTACGCGAGAAATAAC ACTTTTCAAAACT	GGCGAGTTTATTAC GGTGTATGAC	TCTTGCAAT CATTTTAA C	CTTGCAAT CATATTTA AC	0,0	Ots11	41292 800	Ots11	46464 971
Ots_IsoT	T	C	GACTCAGGTAAGGAAA CATCAATGTCA	GAAAGCAAAGCAT TTTATCCACCACTA	AACCAGTA GAATAACC CTAACCCGG	CAGTGGAA TAACC CTAACCTG	0,5, 0	Ots12	10323 252	Ots12	91214 89
Ots_crRA D55400- 59	C	T	CGCAATGAGCCAACCC CT	CTGGTTTGTTCCTG GGCT	A[CA][GA]A C	GA[CA][GA] AC	0,0	Ots12	18236 712	Ots12	17250 308
Ots_1028 67-609	A	G	CTCTGCCATTCATTTGG GCTTTG	GTCTAAAGTGGTCC CCTTGGAT	ACAGAGAG AAGTCCCAG GTG	AGAGAGAA GCCCCAGG TG	0,0	Ots12	21101 745	Ots12	204X XXXX
Ots_1319 06-141	A	T	GGCTCGAACCACCCAG TTTA	TGCCCAACTGGTTT GCAATC	CACGGTTTA CACTCCTAT TA	ACGGTTTA CACTCCAA TTA	0,0	Ots12	21773 920	NA	NA
Ots12_23 066874	A	G	CTCTTTCAGTTGTCTTT GCTCTTG	ATTATGAAAAGGCA TGAACAGGGT	TCCCCACCA AAATTAAGC AAA	TCCCCACC AAGATTAA GCAAA	0,0	Ots12	25387 620	Ots12	23066 874
Ots_pop5 -96	T	C	CTCTTGCTACTTGCAGT GTATCTCA	AGTTTGAGGGCTCT ATTCTGTCATG	TTCTGTTAC TGGAC	CTGTTACTG GGC	0,0	Ots12	29307 827	Ots12	27353 346
Ots_u07- 49.290	G	A	GCTGAGGAAGGATTCT GTATTTGCT	TCGGACAGAGCGC ATCC	CTTTCCCCG TGTTGGT	ACTTTCCCT GTGTTGGT	0,0	Ots12	31470 258	Ots12	30314 171
Ots_cox1 -241	C	T	CACTGAAGTGTAAAGCC ATTGTGATT	GTAAATGTAGTATA CAGTATAGGCATCG TAGGT	CACTACGGT AAGACCAT	CACTACAG TAAGACCA T	0,0	Ots12	31570 301	Ots12	30421 258
Ots_ETIF 1A	A	C	TCTGAAGTACACAAAG GAACACTTG	GAGAGAAAAGGAG AAATGATTGCCATT	CAACTGAA GAAAATAA TATG	CTGAAGAA AAGAATAT G	0,0	Ots12	38462 855	Ots12	34678 104
Ots_GCS H	C	T	GTCTTTTTTAATGATGA CTACAGGTCTTTCAC	GCTACTTTACATAA TACCATTTGAGCTG AGA	TATCTGGGC GGGCTG	CTATCTGG ACGGGCTG	0,0	Ots12	65634 622	Ots12	58052 588
Ots_9689 9-357R	T	A	TCTCCTGAACTAATTT AGACCTCTGAATGT	CCTCATATTGCTTT CATCTGAAGAGAG A	CTGAATGTT TTTTTTAAT CTTT	CTGAATGT TTTTTTTTA TCTTT	0,0	Ots12	67888 781	Ots15	21523 451
Ots_aspat -196	G	C	CCTGAACAGGTACACA CAAACGA	TCCAAGTATGAAT ATGACCAACATGAA T	CACTCTTTA TATCCACAC C[GA]	CAGTCTTTA TATCCACA CC[GA]	0,0	Ots12	68043 508	Ots12	60406 597
Ots_P450	T	A	TGAGCGAGATTTATCA AACTGTCAAAGA	CCCAAGCGGGAGA ACTTACAG	CCCCGAAGT ACTTTT	CCCGAAGA ACTTTT	0,0	Ots12	72832 053	Ots12	65154 384
Ots_brp1 6-64	T	C	ACTCTGGGTCCAGGAG GTTTT	CTGACGAGACCATG CACCAA	AAGTCAGC ATCTTTCA	AGTCAGCG TCTTTCA	0,0	Ots13	42932 54	Ots13	43839 23

Ots_1064 19b-618	G	T	CAAGGGCACATTGGCA GATTTT	ACCGGACCAAAGC ACACA	CAATGATTA ATGATTAAT CCTTC	TGATTAAT GATTCATC CTTC	0,0	Ots13	11736 968	Ots13	11463 845
Ots_HFA BP-34	C	T	CAAGAACACCGAGATC TCCTTCA	TCGGCGGTGGTCTC G	TCGAACTCC GCTCCTAG	TCGAACTC CACTCCTA G	0,0	Ots13	12380 464	NA	NA
Ots_TAP BP	C	T	TTTCTCATCCTTCTCTC TTCCAGTCT	GGACAAACCAGCA CTCCAGAA	CAGCTGTCC AGTTCTG	CAGTTGTC CAGTTCTG	0,0	Ots13	26128 999	Ots30	34770 823
Ots_crRA D27164- 55	A	T	GGAGGCTCTACGTAGG CCT	ACAATATCTGACAC TGACTTGGTCA	AATTTGAAT GACCA	AATTTGTAT GACCA	0,0	Ots13	27591 014	Ots13	25285 990
Ots_myb p-85	C	T	CAAGGGATGTGACAAA TTAATCAAACACATAA	AAGAGGTCTAATAA ATCTCCAATGTAAA AACGT	AGAGCATGT AGTTTTG	AGCATGTA ATTTTTG	0,0	Ots13	31784 059	Ots13	29231 079
Ots_crRA D17527- 58	C	T	TGCCGCTGGATTTATT GACA	GCGTCAGATCAGCT GGTCT	TAGCTCCGA GCTAA	TAGCTCTG AGCTAA	0,0	Ots13	36124 810	Ots11	37324 165
Ots_1124 19-131	A	T	GTGGGTAATCGATGCC AAAGAGAT	TGGCAGTGTTTTCA ACTAGCTTTG	AAGCGACTT GATTATC	AGCGACAT GATTATC	0,0	Ots13	45350 303	Ots32	78532 51
Ots_hnR NPL-533	A	T	TCTTTGATATTGAGCTC ATAAAAGCAAGGT	TCCTTGTTTCATCCA TCAGGCATAAAA	CATTTACCA GTTCTCACA CAC	TTTACCAGT TCACACAC AC	0,0	Ots13	46418 615	Ots13	46705 374
Ots_SL	A	G	AATATTGGCTTTCTGA GAATGCATTTGG	CCAAGATACTTCCT TTAACTTCTCTGTC A	TCAAAGATA TGATTCAAT TAA	AAGATATG GTTCAATT AA	0,0	Ots13	62174 556	NA	NA
Ots_crRA D35313- 66	A	G	TGCAGGAAGAGTTCAG AGAAATCT	GCTCGTTGCAGGTA GAAATGT	TTTAAGATG TAGTT	TTTAAGGT GTAGTT	0,0	Ots13	70298 983	Ots13	65139 272
Ots_U25 67-104	G	A	CATAGTATAGTGATTC GAGTCTGGAGTCT	CGGGCTTTCTTAGG ATATTTTCCTGA	GAGACTGTT GAGAC	GAGACTAT TGAGAC	0,0	Ots13	73139 521	Ots13	68622 537
Ots_IL8R _C8	C	T	CGTGGTGTTGCGCTTC CT	TGTCGGCCATCACT GTCATG	CTGGACGCC GTTACA	TGGACGCC ATTACA	0,5, 0	Ots14	14298 70	NA	NA
Ots_GPD H-338	G	A	CACTAAATATTCCTTA TCATTTCACTAAGT CTGAAGAA	AGCTGATACACAAT CAAAACACAAAAC AT	CCACTACTT AACGTGCTT T	CCACTACTT AACATGCT TT	1,0	Ots14	79965 96	Ots14	11576 466
Ots_CHI 06035945 _4547	C	T	AGCGAGGCTTGCGTTT TACT	GTGCAGTCTGGGCT TGTCCTT	CCGCAACA GATC	CTGCAACA GATC	0,0	Ots14	87879 79	Ots14	12373 492

Ots_U50 49-250	G	T	CAATGTCTAAAGTAAT GGTGGTATTCTTGC	TCTTTGACACACCA TCTGCCAATT	TGGAATGG GTAAGGTGT A	TGGAATGT GTAAGGTG TA	0,0	Ots14	21359 050	Ots14	2257X XXX
Ots_crRA D18492- 65	C	T	GCAGGGCGCAAAGTTC TT	CAGTGAGCGACTGT AATCTGA	TTATGGCTA TTATT	TTATGGTTA TTATT	0,0	Ots14	43620 612	Ots14	424X XXXX
Ots_crRA D57687- 34	T	G	TGCAGGGACGGGGCT	TGCTGTTGTCTTGG GTCTCTC	ACAAATTAA TTAAA	ACAAATGA ATTAAA	0,0, 7	Ots14	3XXX XXXX	Ots14	31889 864
Ots_1040 63-132	C	T	GCGTTACTGGTGTTAT AAACGTTAGC	GTTTATTTAATTAT GAAGGACGATGTTG AAGTCA	CTTTCGTCC TTAGCACAT AG	CTTTCGTCC TTAACACA TAG	0,5, 0	Ots15	27472 51	NA	NA
Ots_NFY B-147	C	T	CAGATGATAGCTTCAG TAAGTGGTTCA	CCGTCCACAGCACA AGACTATAATA	TGTTCCAAT GTAAAATGT ATGC	TTCCAATGT AAAATATA TGC	0,0	Ots15	53130 10	NA	NA
Ots_1247 74-477	T	C	AGTTGTTCTTTTTATAT TGTGTTTTTATTCCATT CCA	GCCAAATAAAAAC AAAGCATGAACAC A	CCACCGCCA TCTGATA	CACCGCCG TCTGATA	0,0	Ots15	10921 418	Ots15	15167 502
Ots_vatf- 251	G	-	CTTTTCGGGTTATTCAT GCTGTTGT	GCAAGCATTTGAAA AACAGACTGGAT	AGACCACA AGATACAGT ACC	AGACCACA AGATAGTA CC	0,0	Ots15	13005 607	Ots17	76640 45
Ots_1122 08-722	C	A	CTGCATGAACGTTAAC TCAAATAAAAGGT	AATGAGTTCTACTG ACATTGTATACTAG AATAAGTATCA	TGTGAGGGC GGTCTT	ATGTGAGG TCGGTCTT	0,0	Ots15	13348 552	Ots15	17637 330
Ots15_18 157381	C	T	TCTCAATGTGATTGAA ATGGATGT	TGTCTGTAGTTTGT GTGTACGGTG	CCCTGGAGA TCT	CTCTGGAG ATCT	0,0	Ots15	13993 075	Ots15	18157 381
Ots_crRA D20887- 70	G	A	CTGCTTGTAGCCGTTT AGC	AGAACACATCTGGC CAGGT	GAAGTCGTC GTTGG	GAAGTCAT CGTTGG	0,0	Ots15	17832 008	Ots20	73780 67
Ots_CHI 06027687 _143477	G	A	GCGAGTGTTAAAAGGG TCAAA	TCTCAAGCCATAAG ACGGGTA	GGAGATAG TCAGGG	GGAAATAG TCAGGG	0,0	Ots15	17999 857	Ots01	34155 766
Ots_1028 01-308	C	A	TGGGACAGAGGTGGG AATTGA	CCCAAAGATGCTTA ACTGAAGATGTG	AGGGACAG TTTCGCAG CG	AAGGGACA GTTTCTCAG ACG	0,0	Ots15	19904 397	Ots15	19971 476
Ots5_709 08626	T	C	TACGGTAGGAAGACTG AATGAGTG	CCCTACCTCTCCAG ATAGCTTGTA	AGCCTCTTC CTCTCTG	AGCCTCTTC CCCTCTG	0,0	Ots15	36201 712	Ots05	70908 626
Ots_1172 59-271	T	G	ACACCCACTTCAACCT CCATAAC	GCCTCAGAGCTTAG CTTGGA	CTCTCCTGA TCACTCTGT	CTCTCCTGA TCCCTCTGT	0,0	Ots16	11947 135	Ots16	11458 685

Ots_1172 42-136	A	G	GTGACAGGAGACAGA AAGAGACATT	TGGTCCTCCCTGTC TCTATCTACTA	CAGCACATA ACTTGACCT C	AGCACATA ACCTGACC TC	0,0	Ots16	29941 952	Ots16	29333 295
Ots_unk5 26	A	G	TCAAGACTGTGCTGTA GTTGTCTAC	CCTCCCCCTTTTCC ACATCAG	CAACATTCC AGTCTGAAA C	CATTCCAG CCTGAAAC	0,0	Ots16	30269 244	Ots16	29699 840
Ots_Myc- 366	T	C	CCTTAGCTGCTCTTTGA AGTTGACT	GGCTATAGAGTGTA TTTACAGCATGCA	TCTCTGCTC ATCTGTC	CTCTGCTCG TCTGTC	0,0	Ots16	30269 853	Ots16	29700 449
Ots_unk1 832-39	C	T	GAAACGTCTATGCTGT CCCCTTTAA	CTGCAGTATTAGCT CTAGTTGAATCCA	CACCACTAG AACTCTC	CACCACTA AAACTCTC	0,0	Ots16	37061 421	NA	NA
Ots17_88 5364	C	A	GTAAAGAAACATGACC TTTTCTGAG	CAGGTTATGGCCAT CATAGTTAAA	TAGCCTTAA GCGCTTCCT GCC	TAGCCTTA AGAGCTTC CTGCC	0,0	Ots17	14052 57	Ots17	88536 4
Ots_1044 15-88	C	T	CCTGAGCATCCCAGTT GAACT	TGTTTTCAATACAC TGCAATTTAGTTTT GGT	TCCTGAAAA ACGACATCC	CTGAAAAA CAACATCC	0,0	Ots17	59936 49	Ots17	55123 67
Ots_SWS 1op-182	T	A	TCAAAGACATCGAACA CAAGAACGA	GCAGGTAAATTCAA ACGTCATCATAAGA A	ATGTACTTT AACGATTCA TTT	ATGTACTTT AACGTTTC ATTT	0,0	Ots17	86561 75	Ots17	73656 88
Ots_crRA D16540- 50	C	T	TGTGTATTCGTCGACC GGA	TCACCTGACCAAAG CACTGG	ATTAAACGT [CA]TGGA	ATTAAATG T[CA]TGGA	0,0	Ots17	11570 673	Ots17	97284 18
Ots_crRA D2806-42	C	A	GCAGGGGCAGACTGA AGG	ACTTCATGCCAATC TCACTAAACA	GTTTGGCAT AAAGT	GTTTGGA TAAAGT	0,0	Ots17	15937 591	Ots17	14222 969
Ots_crRA D22960- 32	C	T	ATCAGGTCTGGGGCGA CA	TTCACCTCTGCCAT CGCC	CGACACCAC TTACA	CGACACTA CTTACA	0,0	Ots17	15937 659	Ots17	14223 038
Ots_IGF- I.1-76	A	T	GGTAGGCCGTCAGTGT AAAATAAGT	GATGGAGGCCACTG TGTTCTTA	CTGCCTAGT TAAATAAA ATA	CTGCCTAG TTAAATTA AATA	0,0	Ots17	17781 778	Ots18	41507 43
Ots18_35 41813	T	C	CCCCAAAAACATCAAG AAGTCTAA	ACATGGTGAGGAA AAGGTAGACTAA	CTACCTACC TTAGTGCTC	CTACCTAC CTCAGTGC TC	0,0	Ots18	45874 5	Ots18	35418 13
Ots18_35 50047	A	G	ATCATCTCTGCTCAGA GGCTATTC	AGAGTGGAAGGAA CGTCTTACACT	TCATTTTTG CAGAGAGA GAAT	TCATTTTTG CGGAGAGA GAAT	0,0	Ots18	46766 1	Ots18	35500 47
Ots_crRA D92420- 25	G	T	AGTGCAGGTCTCCAGA TTTACA	ACCGAAGTGTATGT AAACTTCCGA	CAATCGGA AGTCGG	CAATCGTA AGTCGG	0,0	Ots18	24234 32	Ots18	66951 98

Ots_crRA D33491- 71	C	T	CAGTTCGCTTCTCCAG GGA	TGTGGGTAGCAGAC TGACG	GAGAGCCG AGCTTT	GAGAGCTG AGCTTT	0,0	Ots18	61008 51	Ots18	26993 87
Ots_crRA D12037- 39	A	G	TGCAGGAACTTGCTAT GCT	TGTGGAAAAAGTCA AGGGGTCT	CATTCAAAA AGTAT	CATTCAGA AAGTAT	0,0	Ots18	97703 15	Ots18	10327 070
Ots_crRA D55475- 26	T	G	TGCAGGGTTGGGGACA ATT	AGTCTATTTCCCGA TTTGACTGGA	CCATTTTAA TTCCA	CCATTTGA ATTCCA	0,0	Ots18	17529 243	Ots18	17929 277
Ots18_29 943476	A	G	GTTCATTTTGAAATAA CTGCATCG	CTCTACAAGGTCCA TGCACATTAG	GCCTGACTG GACAACCAT TTG	GCCTGACT GGGCAACC ATTTG	0,0	Ots18	30949 170	Ots18	29943 476
Ots18_30 099101	C	T	CAAATGTAAGGATACG CTTGAATG	GAGTTGCAAGCGAT TACATGTATT	ATTGCATAC TCGAGTCAT CCA	ATTGCATA CTTGAGTC ATCCA	0,0	Ots18	32296 859	Ots18	30099 101
Ots_Hsp9 0a	G	C	GTCGTTTTTCATAGAA AATAGCTCACAGTT	ACAGTATACCGGCT GCCTATTCTATA	ATTTGACTT GTCTTTTT TACAGGAG	ATTTGACTT GTCTTTTT CAGGAGAT	0,0	Ots18	32303 237	Ots18	30105 519
Ots_S7-1	T	C	TGCCATCATAAACAAC CTAACAAGTAACT	CCTGGTTTAAAAAC GGCCAACGTG	ATAAGGTCG CA	AGGGTCGC A	0,0	Ots18	33195 644	Ots18	31651 991
Ots18_32 088284	T	C	CATGAGACACCTGGA GAAAA	ATTTTGATAGTACC TTCTTGGGGC	ATGTTACAT GTA	ACGTTACA TGTA	0,0	Ots18	33781 780	Ots18	32088 284
Ots_pigh- 105	A	-	GCATTACTAAAACTG GTGTGTGGAA	GTTTGGAATGTTTC TCTGATTGTGTAA CAA	TGACCTGAA AATA[TC]AT ATTTTT	ACCTGAAA ATA[TC]ATT TTTTT	0,0	Ots18	38228 583	Ots18	35641 063
Ots_CD5 9-2	G	A	CATGTTACCCAGCTAA AAGTCTATAGCA	TGTTTATCTCTGAG TGAAAAAGGTGTGT	CTAAAATGT CATGTAAAT AT	ACTAAAAT GTCATATA AATAT	0,5, 0	Ots19	95507 12	Ots19	93127 96
Ots_RAG 3	C	T	CATTTCCACGAAAAGC CAGATGAC	ACAGAATAAAGTAT CTTCCTCTTACATC ACTACTAAT	CTCTACAGT ATG	CTCTACAA TATG	0,0	Ots19	36242 326	Ots19	34427 967
Ots_1103 81-164	A	G	CTCTTGTTTGCTATGGG AGATGTAGT	CCGTATCCTAAACC CTTCACTGTT	ATTTGCGTC TTCTCCC	TTGCGTCCT CTCCC	0,0	Ots19	36960 028	Ots19	35228 582
Ots_RAS 1	C	T	TCATAAACATGGTGTC TTTCAGTCAGTT	CTGACATGTGAAAC TACTAAAGCATTTA ATCAC	CAATCTATC ATCGACCAG C	CAATCTAT CATCAACC AGC	0,0	Ots19	37925 628	Ots19	36275 840
Ots_9544 2b-204	A	T	GTCTCTCTCTTTTGCA TCATTACACT	GGACTCTTGAGCTG TCTGGCTATAT	TGGTTCCCC AAATTT	TGATGGTT CCCCTAATT T	0,0	Ots19	40912 881	Ots19	39212 513

Ots19_46 172133	C	T	CACATGGCTCTTTGCT CAAAAT	GCCTACCATTATGT GTTGAATGTT	GCAAATCTC CGATGTAAA GT	GCAAATCT CTGATGTA AAGT	1,0	Ots19	47135 055	Ots19	46172 133
Ots19_46 172427	G	A	CAGTTCCTGACATTCA CCAAAATA	GCAAACAACCCATC ACTAATACAG	TATTCAAAA GGAGCAGTT CAT	TATTCAAAA AGAAGCAG TTCAT	0,8, 0	Ots19	47135 348	Ots19	46172 427
Ots_crRA D20262- 46	A	G	CCTCTGCTGAGTTTGA GGGG	TGAGCAGAGCCTAT GAGGACT	GGTTACA[T C]CCCCAAA	GGTTACG[T C]CCCCAAA	0,0	Ots19	47358 507	Ots19	46397 112
Ots10_21 244146	A	C	CTTCCAGGAGGTATTG TTGGTTAT	TGAACGTAGGTTTG CCATATACAG	CCATTATCA TTAT	CCCTTATCA TTAT	0,0	Ots19	53906 684	Ots10	21244 146
Ots_Est7 40	T	C	GGACTCGTGCTTGAGG AAGATG	TGCATGGCTCCAAC TCCTT	TCTGGATGG AACCGTTAG	CTGGATGG AGCCGTTA G	0,0	Ots20	37510 56	Ots20	39254 576
Ots_crRA D44588- 67	C	T	CGCAAGTCAGCAGGGT GA	TGGGGTTTTAGGCT GGGT	GTGA[AG]CC AATCAAT	GTGA[AG]C TAATCAAT	0,0	Ots20	20333 782	Ots20	25774 788
Ots_Aldo B4-183	T	A	TTTGTGCGTAAAGTCA GGTAGTGT	GTGCATGCCATGAG AACTTTGTTT	CTGTGTGTC TAAGACAAT	CTGTGTGTC TATGACAA T	0,0	Ots21	64771 38	Ots21	52520 21
Ots_AldB 1-122	C	T	GCCATGGAGGACTGGA TGA	GCCACCACTACTTG CTGAGAAAATA	TGTTGGCGA AGTG[GT]GT	TGTTGGTG AAGTG[GT] GT	0,0	Ots21	64801 72	Ots21	52550 56
Ots_1051 32-200	G	T	CGATGTACTGAGGGCA GTGT	GAGTGGAGTTCCTT AATAATCATTGACC TT	CAAGAGTG GCATAAAA	CAAGAGTG GAATAAAA	0,0	Ots21	11799 043	Ots21	95853 79
Ots_1015 54-407	C	G	TGAAAGATATCAATTG TAGTAGTGGTGGTG	ACACGCCAGTCCAC AAGT	ATGGAGGA TTGTGGTTG T	ATGGAGGA TTCTGGTTG T	0,0	Ots21	12050 017	Ots21	97841 11
Ots_GST -375	C	T	CAGCCCGTCCCAAAAT CAAG	CAGGAATATCACTG TTTGCCATTGC	TTTCTTGTA GGCGTCAG AG	TCTTGTAG GCATCAGA G	0,0	Ots21	20087 626	Ots21	18672 650
Ots_GST -207	G	A	GGAGAACATGCATCAC CATTCAAG	TCAGCAAACGAAG GCTATGTAGAAT	ATGAGAGA GTCTTTCTC TGTT	ATGAGAGA GTCTTTTTC TGTT	0,0	Ots21	20087 794	Ots21	18672 818
Ots_redd 1-187	A	G	TTCTGGGTTGCCATAC TCTTTCAAT	AGTTGAGACCTTCA GTTCTTAGGGTAT	ATTCTGACA GCTGTTTTG	CTGACAGC CGTTTTG	0,0	Ots21	27483 610	Ots01	99393 18

Ots_E2-275	A	G	GGTGCCACTTTAGTAT AGCTGCTTA	CCCTACCCCCTGTG TTCCA	CCCCCATAT TGCTG	CCCCACAT TGCTG	0,0	Ots22	64192 64	Ots22	54570 12
Ots_123048-521	A	C	CTCAACAGTGCACCTC CCTTAATT	CCAAACACACCCTT CCATAATCTCT	TCACATCCA ACTCAGTACT	CATCCAAC GCAGTACT	0,0	Ots22	94784 78	Ots22	80333 23
Ots_parp3-286	A	G	AGTCAGTGTGGTGTA GTGAAGAGA	CATTTGTGGAGTGT TTATTGAACAGTAA CA	AGTTACAAG TGGTGTTTC A	ACAAGTGG CGTTTCA	0,0	Ots22	10728 055	Ots22	95051 20
Ots_u07-17.135	A	G	CTCGCCTCTGTCATTGT ATTACCTT	TGACACACGAGCCA TTTTGATGAT	AAAATGTAC CACATACTT GT	AAATGTAC CACATACT CGT	0,0	Ots22	12134 293	Ots22	11164 448
Ots_CD63	A	C	TGCATGTTTTCTAACTG TGTTTTTGTT	TGAATGCCCCCAT CAACA	AGATCATGG GAATCATAT	ATCATGGG CATCATAT	0,0	Ots22	14629 707	Ots22	13378 891
Ots_110551-64	C	A	GAGTGGTCAAGGTTTC AGTTTCTG	GAAATGGACAGAC ACAAGGTCAAAC	ACGCTCGGA ACATT	ACGCTCTG AACATT	0,0	Ots22	15838 315	Ots22	14420 914
Ots_u1002-75	T	C	CCGCCTTTCCACCTTC TC	TCAAACGAGAACA CACTAAGGTTGT	ATGGCCCTT ACACTATC	TGGCCCTT ACGCTATC	0,0	Ots22	16981 803	Ots22	15468 014
Ots_tpx2-125	C	T	TGTTGTAATCTTTCTGA ATATTTGCTTGCTT	TCTTCCAAATTGAG CACAAAAGCAT	CAGGCGGTT CTCC	CAGGCAGT TCTCC	0,0	Ots22	21879 673	Ots22	20362 912
Ots22_32650802	G	A	AAGGAGCAGGAGATG TTATTGAAG	ACTAGGTACTTCAC TGGTCCACTG	GGGAGAGG AGGCCTGTC TTTA	GGGAGAGG AGACCTGT CTTTA	0,0	Ots22	34412 033	Ots22	32650 802
Ots_CHI06105101_16717	C	T	AAGGCCGTGAACATCT GTG	ATCGCAGGCTAGCT TTTCAA	CCTCACATA CTCCCTT	CCTCATAT ACTCCCTT	0,0	Ots23	18000 122	Ots23	51243 25
Ots_CHI06105101_18523	A	G	GCGGTGGGATACCTCC TCTA	GCGAGAAAAGCAC TGAATGA	GGCGGCTCG GAAAATTAT TTT	GGCGGCTC GGGAAATT ATTTT	0,0	Ots23	18003 825	NA	NA
Ots_106747-239	C	A	ATCGAGGATGCCTCAA AGACATC	GTTAGACCCACCAC CAGTCATC	CCC GCGGTG AGTAT	CCC GCTGT GAGTAT	0,0	Ots24	95681 30	Ots24	17321 046
Ots_PEMT	C	T	AGAGCATTC AATTTAA AAGCTGAAAACGA	CTTTGATCCCTGCT TGCAGTATTTT	TGC[AT]TTG CTAAGACTT G	TGC[AT]TTG TTAAGACT TG	0,0	Ots24	10785 871	Ots24	15921 271
Ots_117370-471	G	T	GTTGGCTCCTTCAATTC AATTTGGA	TGCAAACACAGAG GAAAGGGATTT	ACGGAACA AATAAGAC ATTT	CGGAACAA ATAAGCCA TTT	0,0	Ots24	12157 474	Ots24	14581 248
Ots_107074-284	A	T	CCCACTTCCAGAGCCT GAA	TTTTCCATGGCTGT GTGTA CTGT	ACCGTAGCT GCACCTG	CGTAGCAG CACCTG	0,0	Ots25	81428 48	Ots11	16750 220

Ots_GTH 2B-550	C	G	CACAGGAAGGACGTGT TTTGATG	TGACTACCCGTTGT ACCAATGAAC	ATAACATCT GCAGCATTAA	ATAACATG TGCAGCAT TAA	0,0	Ots25	26054 245	Ots25	27327 552
Ots_myo 1a-384	A	C	CTCCCCCTGGACTTT GG	GCTCTATTGCACCG TGTTCTG	ACAGATCCA TCCACCACT	AGATCCAG CCACCACT CACATCAC	0,0	Ots26	10237 957	Ots03	39283 854
Ots_1128 76-371	C	A	GCCTACAGCAAATTCA GCTACACAT	TGGACCTTCAATCA TCACAGCTT	CATCACAAC GATGTGTG	AACTATGT GTG	0,0	Ots26	30XX XXXX	Ots26	29108 047
Ots17_22 360456	T	G	ATGGTTAAATTGACTC CTCCCTAT	GCCTACTACTGTTC TGTCATCTGC	AGTCTGTCTG TTGT	AGGCTGTC GTTGT	0,0	Ots27	14067 908	Ots17	22360 456
Ots_RAD 4543-52	T	C	TCTTTGGACTGTGTAT ACCAGGTGTA	GCCAGATGCTGTGT GTGTTT	TACATATGA CTAATGAAA	TACATACG ACTAATGA AA	0,0	Ots27	17341 235	Ots27	12866 217
Ots_Endo RB1-486	G	A	CCTTTGGGTCTGCTTG AGGTT	GGAGCCAAATCCTA ATGCTGAAGTA	TCCTTCTCA CGCTTCT	CTCCTTCTC ATGCTTCT	0,0	Ots28	93101 15	Ots28	81874 90
Ots_1088 20-336	G	A	TGAAATAAATTGTTCT GTTGATATGTGAATTT TGGA	CAACGACACACCA ACAACGT	ATTGCCCAT CTCAGAATA	AATTGCCC ATCTTAGA ATA	0,0	Ots28	12014 613	Ots28	10815 622
Ots28_11 023212	A	G	AGAAAGCCATCATCAT GAGACC	ACAAACAAACAAA AATGGTCAGAA	AACGTGAC ACAAT	AACGTGAC ACGAT	0,0	Ots28	12231 157	Ots28	11023 212
Ots28_11 025336	A	C	TGCAATATAGAACAAA TCCGAAAA	AATAACCTTGGCT TCACATACAT	CAATGAAGT TAATTTAAT TGG	CAATGAAG TTCATTTAA TTGG	0,0	Ots28	12233 225	Ots28	11025 336
Ots28_11 033282	G	A	GGCTTTCTGATGATCTT GAACCTT	AGTGTGAGAGAGA GGAAGTCCCTA	TAAAAATG GTTGATATG TA	TAAAAATG ATTGATAT GTA	0,0	Ots28	12241 662	Ots28	11033 282
Ots28_11 062192	C	G	AGATGATATGGATTG CTGTGTGT	TTGAACATAACGAT CAGAGAAAGA	TTCTCAAGT CCTACTCAA CTG	TTCTCAAGT CGTACTCA ACTG	0,0	Ots28	12266 158	Ots28	11062 192
Ots28_11 070757	A	G	TTTTGGAACCCTTTTAA CTACGAG	ACATCAGTATAGCA GAGGAGAGGG	ACCCATGAA TAAGGACG AGAG	ACCCATGA ATGAGGAC GAGAG	0,0	Ots28	12274 804	Ots28	11070 757
Ots28_11 071377	T	C	ATTTGCTGTGTGTGGA GTGAAT	GTAGTGACAGATGC TCTTGGAGG	CATCTTAGC CTCTCTGAC CCC	CATCTTAG CCCCTCTG ACCCC	0,0	Ots28	12275 402	Ots28	11071 377
Ots28_11 072994	C	T	GGGAGACTTAAAAACAA CCTCAAAA	ACCTGCAACCTTCT ATTCAACAGT	CCATATGTC GCTTGT	CCATATGT CGTTTGT	0,0	Ots28	12277 000	Ots28	11072 994
Ots28_11 073102	T	A	GGTGAGCCATTTCATAA CAATCTT	TGTTATCCTGGATC ATTCAAGAGA	ACATTACTT TTCAAAAAT ATT	ACATTACTT TACAAAAA TATT	0,0	Ots28	12277 108	Ots28	11073 102

Ots28_11 073668	T	A	CCTAAGAGGAGACGA GCATTACAG	GGTAAATCAACATA TGACCACTCG	TACAGTTTC CTGTCTGA	TACAGTTTC CAGTCTGA	0,0	Ots28	12277 674	Ots28	11073 668
Ots28_11 075348	G	A	CATTTCAAAATTAGGA GGTTAGGG	AGATGAGAGCTGTG GCCTGT	GTGTGAAA GGGGAGAA GGGCT	GTGTGAAA GGAGAGAA GGGCT	0,0	Ots28	12279 292	Ots28	11075 348
Ots28_11 075712	C	T	GCTTAAACAGCTGCTA TTAGGACA	TAAGGATTTGTTGC CAGCTCTAAT	GAAAACTCT GCCCTG	GAAAACTC TGTCCTG	0,0	Ots28	12279 656	Ots28	11075 712
Ots28_11 077016	C	T	AAAATATGTGCAACAT CCAATGTC	ACACAAGCTGGCTG AAGCTAAT	GTCAAACCA ACTTTGCCA AGG	GTCAAACC AATTTTGCC AAGG	0,0	Ots28	12280 918	Ots28	11076 976
Ots28_11 077172	G	A	GTTTTGCCAGAGAGAA TGTACAAA	TAGTGGTTAGAGCA TTGGACTAGC	ACACACAC AAGAGACA CCCAC	ACACACAC AAAAGACA CCCAC	0,0 4	Ots28	12281 112	Ots28	11077 172
Ots28_11 077576	A	G	TGTGCGGAATTACTGA TAATTGAC	GCTCTGCATTTTAC AACACTGCT	GAAGGCCA AATAAAATT G	GAAGGCCG AATAAAAT TG	0,0	Ots28	12281 512	Ots28	11077 576
Ots28_11 095755	A	T	CCAATGGTGATTTTAG AACCATTAC	AAAACAGAGTATG GATCAACAGCA	AGAGTTGA ATGGC	AGTGTGTA ATGGC	0,0	Ots28	12299 996	Ots28	11095 755
Ots28_11 143508	G	A	ACCTTTTAGCCAGTGA CAACATTT	ATGCAAGAACTCT CGACGATAG	TTCACGTAC GGCCCAT	TTCACATA CGGCCCAT	0,0	Ots28	12341 541	Ots28	11143 508
Ots28_11 160599	G	T	GTGCATATTTTACGTG GTTGAAGT	ATTCCATTTACCC ATATGAATTT	CTCTCTGCT TGC GTT	CTCTCTGCT TTC GTT	0,0	Ots28	12359 222	Ots28	11160 599
Ots28_11 164637	C	A	TGATTTGACTTTTTGTG GTGTTTT	GTTCCAATCTGTTT TTGCTCTCTT	CTGGCGGG GTCTGGG	CTGGCGGG GTATGGG	0,0	Ots28	12363 212	Ots28	11164 637
Ots28_11 186543	A	T	GGCTTGCCTTTAGATA GAATCTTG	AAATCTCACAAGTC CAAAAACAAA	AAAGCTGAT TAAAAA	AAAGCTGA TTTAAAA	0,0	Ots28	12385 919	Ots28	11186 543
Ots28_11 201129	T	G	TGCGAGATTTATCTAC TTGTCCAG	GGTAGTTTTGTACG CAATTGCTAA	ACTGAAGG AATTTAAC	ACTGAAGG AAGTTAAC	0,0	Ots28	12400 459	Ots28	11201 129
Ots28_11 202190	T	C	GCTAAATGTAAATCGA GTGGCTGT	TACATGGGTCCTCT CAGTGTTCTA	CAAAAGTCT GTATTTTCA AAA	CAAAAGTC TGCATTTTC AAAA	0,0	Ots28	12401 520	Ots28	11202 190
Ots28_11 202400	C	T	CCCTCCAAAAAGAAAA CATTTGAT	AAATTGGCTAATCA AACACTGGTT	GACACACTC ACGA	GACACACT CATGA	0,0	Ots28	12401 729	Ots28	11202 400
Ots28_11 202863	C	A	GAGGATGGATGAGACT TTTCAGAT	GCTCTTTACCGGGT TTATATGAAG	ATAAAAAA TTCTGCGTG AATG	ATAAAAAA TTATGCGT GAATG	0,0	Ots28	12402 193	Ots28	11202 863

Ots28_11 205423	A	G	TTAAATCACCCAGAGC TTGTTAGA	ACCTGACCTAGATA ACAACCACAA	CCTGCACAC ATGTCAAAC CG	CCTGCACA CGTGTCAA ACCG	0,0	Ots28	12404 734	Ots28	11205 423
Ots28_11 205993	C	T	GCTGCTATTTCCGACC TTACAATA	ATCAAGACAAAAC ACTCACCAGAA	GCTATTAAA AGG	GTTATTAA AAGG	0,0	Ots28	12405 298	Ots28	11205 993
Ots28_11 206740	T	C	ACTTTGAGGACTTACT CCTGTCCT	CTGGAGAAAGACA AGATGATGATC	CCTTCCCTC CTAGGGCA ACGT	CCTTCCCTC CCAGGGCA ACGT	0,0	Ots28	12406 045	Ots28	11206 740
Ots28_11 207428	T	G	TATACCTTTGTAGCAT CCCTCTCC	CATATAAAGTGGAC AGCGTTTGAC	GTTGGGAGC GTCCCAAAA TGG	GTTGGGAG CGGCCCAA AATGG	0,5, 0	Ots28	12406 663	Ots28	11207 428
Ots_trnau 1ap-86	G	T	GGACAAGTTGAAACAG ATCAGGAAGT	GCCACTGGATACCA TCACTTCAAA	AATCCCTCC TTTTTCC	TCCCTCATT TTCC	0,0	Ots28	18849 666	Ots28	17646 942
Ots_OTS TF1- SNP1	G	T	CGGACAAAGAGCTACA GAAATGC	CGTCCCTCTTCACG CATGA	CCGCCACCT TGGCT	CGCCACAT TGGCT	0,0	Ots28	27509 692	Ots28	25845 386
Ots_1224 14-56	C	T	GCACCGTATCAACGAG CTCAT	TGCATGGATTTCCT TTGTGTTGTTG	TGTATGACC TCTGACCTG T	TGTATGAC CTCTAACCT GT	0,0	Ots28	28489 780	Ots28	26647 710
Ots_CCR 7	C	T	CTGCTCACCTGCATCA GTGT	CCATGGTGGTCTGG ACGAT	CCACGTAGC GATCG	ACCACATA GCGATCG	0,0	Ots28	37080 463	Ots28	34762 262
Ots_9766 0-56	A	T	TTCCCTAATCTGACGT ACTACCAACT	CGCCACTGACGTTC ATTCCA	ACGAGACA GATATTC	ACGAGACT GATATTC	0,0	Ots28	37155 946	Ots28	34830 271
Ots_crRA D18289- 33	T	C	GCAGGGAAAACTGGTC AGGA	AGGTGAACCTCCGT CCCA	GAATGGTGT TAAAT	GAATGGCG TTAAAT	0,0	Ots29	78932 96	Ots34	89707 78
Ots29_23 344676	T	C	GAACTATCCTGACTCC CATTGAAA	CTGAGTTCCTCCTG GTTGTTATG	TGCAAGTCC TTCAAAGGC TCA	TGCAAGTC CTCCAAAG GCTCA	0,0	Ots29	12582 492	Ots29	23344 676
Ots_1110 84b-619	C	A	TTGTGGAATTACACCT TCAGAGTTCAAT	GCCTGTTTGGCTTT CTTAAACTGAT	TCCATGG[A T]AACGGAC AAT	TCCATGG[A T]AACTGAC AAT	0,0	Ots29	16022 207	Ots29	19749 885
Ots29_18 791740	T	G	GTTTTGGTGTGGTCTC AAATCC	CGGCACCTGGAAAC AGTC	CCTATGAAG TT	CCGATGAA GTT	0,0	Ots29	17054 573	Ots29	18791 748
Ots_1083 90-329	G	C	GAGGTTTGTACTGTC ACCCATAGA	CCTGCTGTAGCAAA CTGTCTCAAA	CTACTTATG TAGCATTTT AA	CTACTTATG TAGGATTTT AA	0,0	Ots29	19105 883	Ots29	16533 976

Ots_9622 2-525	C	T	GCTCTTGCCCATCTGT AGGAT	GGCGCAACATATGT ATTAAGCAACT	TGTAGCTAA TTTTAAGTT CTC	AGCTAATT TTAAATTCT C	0,0	Ots29	23142 347	Ots29	12489 280
Ots_U24 46-123	C	A	CTGGTCTGTGACGTCA AAATGATG	AGCTAGACCAGGCC ATTTGAG	CTGCAACTC GACGCAAG	ACTGCAAC TCTACGCA AG	0,0	Ots29	23877 989	Ots29	11728 129
Ots_1174 32-409	A	G	TCATCAAAACATGCCT CTTCTGTGT	TGTTGAACCTGTCA CTCTGTCTTC	TTTAGACTT TGCTCTATA ACAG	ACTTTGCTC CATAACAG	0,0	Ots29	25338 465	Ots29	10328 032
Ots_crRA D255-59	T	C	TGCAGGAGCTGTGATG GG	GTACGGAGCGTCAC TGCT	AACTGTTCA AACCC	AACTGTCC AAACCC	0,0	Ots30	69759 04	Ots30	40093 591
Ots_RFC 2-558	A	-	AAGGTCTACTCCGGTT GTATTCGGT	CAATACGACAGTAC CGGTGTTAAACT	TGCATGTAA CAAATAAC AT	TGCATGTA ACATAACA T	0,0	Ots30	17770 066	Ots07	27438 208
Ots_PGK -54	T	A	CTCATACTTTGTACCTG TGTGTTCCA	CGACCCAAGTGGCT CATCAG	CCACCATCA AGCACTG	CCACCATC ATGCACTG	0,0	Ots30	19962 166	Ots30	23782 920
Ots_1307 20-99	A	G	CGGTCATTGTAAATGT CAACGGTTT	TGCTTGCATGTTCT TGGTGTAGTAA	CCTGTCTCA TTCCC	CTGTCCCAT TCCC	0,0	Ots30	24569 258	Ots30	17928 099
Ots30_17 330688	T	C	CTGACAAAAGTGATCT GCCTGA	TGCTTGGTTACACA GTTTGACA	TGTGTCTGA GA	TGTGTCCG AGA	0,0	Ots30	25291 039	Ots30	17330 706
Ots30_17 330452	G	C	CACAAATGTGACCGTT TTCATC	TTGAACCAGGGTGT CTGTAGTG	CATGTCAGT GC	CATGTCAC TGC	0,0	Ots30	25291 293	Ots30	17330 452
Ots_EP- 529	A	G	GCCCTGCCTGCAACTT C	GAAACCAACGTCTT GATGTAGACCTA	CAGTGTCAT TTTCGGC	ATCAGTGT CATCTTCG GC	0,0	Ots30	26852 592	NA	NA
Ots_crRA D25367- 50	T	G	ACTGCAGGCGTCATGC TT	TGGACAAAAGACC ACAGGCT	GTATATTTA GAATG	GTATATGT AGAATG	0,0	Ots30	34187 799	Ots30	80425 26
Ots_u07- 18.378	A	T	GGAAACCAGCTAGGAT TCAGGAA	CGTTATATGGTTTG CTTGTTTGCATA	ATATGGTAT GTAGAGGCT AGTTA	TATGTAGA GGCAAGTT A	0,0	Ots30	36420 567	Ots30	61306 60
Ots_U23 62-227	A	T	TCGTGGATTGTGGCTT ACGT	GGGTGTTTAAACAAG TAGTCCCTTCA	AAGAAGCA TTTTTT[GT][GT]	AAGAAGCA TTTATTTT	0,0	Ots30	38784 785	Ots06	14700 070
Ots_U23 62-330	A	G	AATGGGTAACAAAGA AATAGCTAGCTACTT	GACAGACCACAGT GAAGGTGAAA	ACTGGGAA GATTGTTTG	CTGGGAAG ACTGTTTG	0,0	Ots30	38784 886	Ots06	14699 971

Ots_crRA D57520- 66	T	G	ACAGAGCTGTGTCTAC CAGA	ACCCTCTCTTGGCC TTGC	TTTTTGTTT AAAAG	TTTTTGGTC AAAAG	0,0	Ots30	39954 606	Ots30	13323 75
Ots_P53	G	A	GGAACCTCCTCTCCCG TTCTG	GCACACACACGCAC CTCAA	CTGGGTCGG CGCT	TGGGTCGA CGCTC	0,0	Ots30	40916 627	Ots30	24175 91
Ots_1096 93-392	T	G	TCTCCCTCATTCCCATG TCATATCA	GGGAACGTATCAG GTGAGTGT	TCCGTTAGT TCATCCTGG	TCCGTTAGT TCCTCCTGG	0,0	Ots31	93693 78	Ots13	74025 50
Ots_unk1 104-38	C	T	TAACCATGACTTCTAT CAATCACCCC	CCTCCATACATCGT CAAAGCTGTA	CCACTAAGG ATTACGTTA CG	CACTAAGG ATTACATT ACG	0,0	Ots31	14800 538	Ots31	21291 286
Ots_ntl- 255	T	A	TGCAGTTACAAGCCTA AGACAATCT	CAACTAAAGTAACA CACCAGCAACTG	ATTCTTCCT C[TC]ACAAT TG	ATACTTCCT C[TC]ACAA TTG	0,0	Ots31	20616 326	Ots31	15370 000
Ots_1011 19-381	T	C	TTTTCTAGGACAGGTT GCTTGCA	CCAGGTTTCTTTAG CCTACTTATTCTTTA CA	TGCCACATG ATAATTGA	CCACATGG TAATTGA	0,0	Ots31	21772 790	Ots02	58361 347
Ots_txnlp -321	T	C	CCTTCAAACCTAACACA TCATAGACATGCTT	TTATCAAACCTGAAG GCGGATTTACTGA	TCTGGCGGA TTTACA	CTGGCGGG TTTACA	0,0	Ots31	23141 496	Ots31	13146 728
Ots_1132 42-216	C	T	GAGGCCTAATGTCTCT TGTGACT	GACATCTTCAACAA GTGTTCAATCACC	ATTACCAAC GGAGAACC	TTACCAAC AGAGAACC	0,0	Ots31	31397 037	NA	NA
Ots_sept9 -78	G	A	GTCGATTACCGTTAGC TTCATCCT	ATTCTCTCTGTGTCT CTCTCTGTCT	CTCTTCGAT GTCTAGACA	CTCTTCAAT GTCTAGAC A	0,0	Ots32	55431 76	Ots32	93903 57
Ots_DDX 5-171	C	T	ATGACCAATTGAAGAG TTCTTCCGT	CAAAGCCAAACGTC ACATTTACACT	TTCATAATT GAACGATTT CA	CATAATTG AACAATTT CA	0,0	Ots32	71833 41	Ots32	59485 80
Ots_1181 75-479	C	T	TGCGCGTCTCATTCAA CCAT	ACCTTACGTCCTAG GTAGGAAACA	AGAATGAA GTGAAAAG AA	AGAATGAA GTAAAAAG AA	0,0	Ots32	76315 74	Ots32	54361 60
Ots_1298 70-55	A	T	GCATGTAACACATTAT TTGGCATATGTACT	CAGTACACTGGAGA TTTGCAATGTT	ATGCATTCA CCTGTATTA T	TGCATTCA CCAGTATT AT	0,0	Ots32	10246 920	Ots32	29798 52
Ots_Ostm 1	C	G	CCAGCCCCGTAACACA CAT	GAGAGGAAGCAGA AAGGTCGTTTAA	CCGTGGTAT TGTTTCAA	CCGTGGTA TTCTTTCAA	0,0	Ots33	27432 27	Ots33	45699 55
Ots_NA ML12- SNP1	A	G	TGCCACCTCAGTTTTA GTGTTATATCC	AGCGCCAACCTGTC ACT	AAACCATTT TCATTCTTT TG	CCATTTTCA CTCTTTTG	0,0	Ots33	85237 57	Ots33	97778 02
Ots_CRB 211	A	C	CAACGCGGGAATGGCT TTTAA	GCCAGAGTCGCCAA AATAGTAGAAT	CTACCGTAC TGAATCTC	CCGTACGG AACTC	0,0	Ots33	28420 240	Ots33	29785 703

Ots_TGF B	C	T	GCCTCACATTTTACTG ATGTCACTTC	GAGCAGATCTCTTC AGTAGTGGTTT	AGCCTAGCT CTCGGAAG	AGCCTAGT TCTCGGAAG	0,0	Ots33	28445 695	Ots33	29760 263
Ots_1064 99-70	C	G	ACTCTATCATCGGCAG GACCAT	ACCGTAAGTGTGGT TGTGTTTCATTA	CATTTTTCA GAATTGTAT TC	CATTTTTCA GAATTCTA TTC	0,0	Ots33	37411 470	NA	NA
Ots_TLR 3	C	T	TGCACCTGCGAGAGCA T	CTGGCGTTTGTTC GTTTCAG	CTGTGGTTT GTGGCGTG	CTGTGGTTT GTAGCGTG	0,0	Ots34	72341 89	Ots34	73652 14
Ots_crRA D24807- 74	A	T	TGCAGGAGAGCAGGGT AGA	CGTGCCTAACATCA TGTGCA	ATGATAAT	ATGATATT	0,8, 0	Ots4	44035 017	Ots04	36172 590
Ots_crRA D36152- 44	C	T	CAAAGTGCAGGTGCTG GC	CCAGCCAGGTGTTG AGCA	CTGCCACCC TTTGA	CTGCCATC CTTTGA	0,0	NA	NA	Ots12	13203 734
Ots_afmi d-196	G	C	CGTGGAGTAGGTGGTT ACAGTTTAT	CTCGTAACAAGCTA CTGTAGTGTACT	CAAAGTCA AAGATCCTA TAAA	AAGTCAAA GATCGTAT TAAA	2,0	NA	NA	Ots32	17096 334
Ots_1266 19-400	C	T	GGATGGTTGTCATTTT TCTGCAAA	CCGGGATACAATAA TAATATTTGGTTAA GAGTTTTTT	AGAAAGTTC TAGAAATA ATT	AAAGTTCT AGGAATAA TT	2,0	NA	NA	NA	NA
Ots_Est1 363	A	T	GGTGATTTTGCCACAG AGTAGAGAT	AGTGTTAAATGTAA CTTGCATATACAGG CAAT	CCATCCTGT CTTGTCTG	CATCCTGTC ATGTCTG	0,0	NA	NA	Ots03	82923 51
Ots_U23 05-63	T	-	TGTCATCTCTATTGCA ATCTCAGTAGATTTCT AT	CCAGGTCGTCTTTA TTGCAGATTATCA	AATGTCATA TAGAAATCT AC	AATGTCAT AGAAATCT ACTG	2,0	NA	NA	Ots14	398X XXX
Ots_u07- 07.161	C	T	GTCAACAAATGCAGGT AACATAAATGGT	GATGCAAACACCTG TGAAATTGTGA	ATCAGTGAC ATAAGTTGT CCA	TCAGTGAC ATAAATTG TCCA	0,0	NA	NA	Ots04	37799 853
Ots_u202 -161	T	A	CACTTTTGACTTTACAT GGAACCTAACTCAT	GGGACTTCACTTTC TACAAACATGTCA	AGCTAGTGC TTAGCAGCT A[AC]	AGCTAGTG CATAGCAG CTA[AC]	1,5, 0	NA	NA	NA	NA
Ots_Cath _D141	T	C	CACTTGTTCTGCACAC TACTTGTC	CACACATGGATTTT GCCTGTCTAAA	TGGGAAGC AATCAA	AATTGGGA AGCAGTCA A	0,2	NA	NA	NA	NA
Ots_1104 95-380	G	C	GCCTAGGTATGTACGA AACTTCACA	AGGCTTTTTTCAGAT GGTCGTATGA	CATAGAC[A G]GGGGCCA T	CATACAC[A G]GGGGCC AT	1,9, 0	NA	NA	NA	NA
Ots18_34 17174	A	C	TGAGGTATTACTTGCT GAGTTTGC	CAAGAAGATGTGA ACTAATTCCCA	CTGAATCCT GTAAG	CTGCATCCT GTAAG	0,0	NA	NA	Ots18	34171 74

Ots_1239 21-111	A	G	TCGCTAGGCAGAAATA TAGGGTTCT	GAGCATGGCGCTTG CA	TGCTAAATG GCATATATT AT	CTAAATGG CACATATT AT	0,0	NA	NA	Ots18	25247 852
Ots_unk7 936-50	C	G	ATGGGTTGGGATTATG GTTCAATTGT	CAAAATGGTTACTT GCATAGTCTTTTGT	AGACATGTA GCTATGTAG GTAA	AGACATGT AGCTATCT AGGTAA	0,0	NA	NA	NA	NA
Ots_9955 0-204	C	T	TGACAGATTTTCACCTT TAACTAGCTAAGC	GCAACCTCTTTCAC ACTTCAGTAAC	AAGGCTTTG GTTGTTTG	AAGGCTTT GATTGTTTG	0,0	NA	NA	NA	NA
Ots_crRA D60614- 46	G	T	TGCCGTGAGAACTGG TCA	TTTCCTCCTCTCTGC CTCA	AAGATGGT ATGTAT	AAGATGTT ATGTAT	0,0	NA	NA	Ots12	72156 356
Ots_OTD ESMIN1 9-SNP1	C	A	GGTCTGTCTGTCTGTCT ATCTGTCAATG	TGTGTGTCTTTGTTC ATTCCTACCA	CCAGTCATG GGTCATT	TCCAGTCA TTGGTCATT	0,0	NA	NA	NA	NA
Ots_TCT A-58	C	T	ACCAGTACCTAAACGT TAGAAAGCAA	CGTTAGTTAGCTAT GTCTGAAAGGCA	CTGCCATGA AGTGCTAG	TGCCATGA AATGCTAG	0,0	NA	NA	Ots02	89930 26
Ots_MH C2	T	G	GTCCTCAGCTGGGTCA AGAG	GTAGTGGAGAGCA GCGTTAGG	CTGGAGCGT TTCTGTA	CTGGAGCG TGTCTGTA	0,5, 0	NA	NA	NA	NA
Ots_1100 64-383	C	T	AACAAAGAATGTTAAA CACCAAACAGGAA	GTGCAAGGGACCTA GCTAATCC	CTACGTAAT GAACGTTAG CT	ACGTAATG AACATTAG CT	0,0	NA	NA	Ots27	81410 61
Ots_U21 2-158	G	A	CCCCATATGAGACGCT ACAGTAATG	CAAATGCCCTCTAA GCAGACCTT	CTGGAAGA AGGCCTC	CTGGAAAA AGGCCTC	1,5, 0	NA	NA	Ots27	14522 192
Ots33_19 359879	T	C	AGCGCCTGTTTTACAT AAACACTT	GTGAGTACCGTAAA GACTGAGCAA	AAATAAAC GCTGGGTCT AATT	AAATAAAC GCCGGGTC TAATT	0,0	NA	NA	Ots33	19359 879
Ots2_382 64269	A	C	GTATGAGTTGTGTGGT TGCAATGT	CTCTAGCCTATTGC ACAATGTCC	TCCCTTGTC TATGGTATA TCT	TCCCTTGTC TCTGGTAT ATCT	0,0	NA	NA	Ots02	38264 269
Ots18_34 26299	T	A	TTATTTTGGGCTTCATA TGGTTCT	GGTCATGATGTTGA TATTTTGGGA	AATGCCATT TTGT	AAAGCCAT TTTGT	0,0	NA	NA	Ots18	34262 99
Ots_U51 21-34	A	G	CCAGAGGTTAGATGGC CCTTT	CTGAGCCAGAACCA CAAATTGAATT	AGGGTCTCA TGCTCCCT	AGGGTCTC GTGCTCCCT	0,0	NA	NA	Ots05	20893 356
Ots_crRA D60620- 51	A	G	CAGGCAGTCACTGAGT CCG	TTTGAGCACCGTTT CCGA	GTACGGAA AAAACA	GTACGGGA AAAACA	0,0	NA	NA	Ots21	24004 835
Ots_ARN T	G	T	CCACTGGCTGTGGAGC TT	GGGTTTCAGTGATAG TTGGGCAAAT	TACAGATGT CATTTTAC	CTACAGAT GTAATTTTA C	2,2, 0	NA	NA	Ots23	24857 375

Ots_GH2	A	T	GCGTACTGAGCCTGGA TGACA	CCCCCAGGTTCTGG TAGTAGTTC	TGACTCTCA GCA[TA]CTG	TGACTCTCT GCA[TA]CT G	1,8, 0	NA	NA	NA	NA
Ots_IL11	T	C	CCTCCAGATGAGACCC ACTCT	CAAAATGGTGCTCA AACGACTTCA	AGCTCCATG CGGACT	AGCTCCAC GCGGACT	0,0	NA	NA	Ots20	20625 334
Ots_1182 05-61	T	C	CCATACAGCCAGTCCA GGTG	ACTGGACAGGGCTG GGT	TAGTAGCCC CTACACCTC	TAGCCCCT GCACCTC	0,0, 4	NA	NA	Ots33	32594 959
Ots_crRA D26081- 28	T	G	GGGAGAGGGAGACGT GGA	TCACCAGCTCCTCC TCCTC	TGGAGGTG GAGGAG	TGGAGGGG GAGGAG	0,0	NA	NA	NA	NA
Ots28_11 210919	C	T	AGTGCTCCATGCTGGA GTTT	GATGAAGCAGAAG GAGAGGCT	GACCTCAAG CAGTCAG	GACCTTAA GCAGTCAG	0,0	NA	NA	Ots28	11210 919

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Section 2: Genetic Baseline Expansion

Introduction

Distinct population aggregates of Chinook Salmon (*Oncorhynchus tshawytscha*), steelhead trout (*O. mykiss*), and the species *O. nerka* (Sockeye Salmon and kokanee), have evolved through the cumulative effects of selection and genetic drift (Waples 1991). The homing behavior (philopatry) displayed by Pacific Salmon means that fish typically return to spawn in their natal rearing sites or stream of origin. This distinctive life history attribute can significantly restrict gene flow, shape regional variation, and influence demographics among naturally reproducing populations (Hasler and Scholz 1983; McIssac and Quinn 1988; Quinn *et al.* 1991). Genetic differentiation is most easily resolved among populations that are geographically distant, where degree of gene flow is generally correlated with relative migration distances and adjacency in stream networks. However, local adaptations and the distribution of suitable spawning habitat within stream networks may influence finer (regional) scale genetic structure among watersheds in close proximity (Beacham *et al.* 2006; Matala *et al.* 2012). The natural phenomenon of immigration or straying (a homing miscue) buffers the loss of genetic diversity in salmon populations (Milner and Bailey 1989), but the rate of straying exhibited by wild fish is generally low (Quinn 1993; Heard *et al.* 1995) and genetic structure between populations may persist despite moderate gene flow from straying (e.g., Neville *et al.* 2007). Some evidence indicates that hatchery-origin fish exhibit a higher rate of straying which may be affected by changes in fish passage protocols, transport through the hydro system, artificial rearing practices, or inadequate acclimation (imprinting to natal waters by juvenile salmon). An elevated rate of immigration between populations may erode local adaptations, and lead to changes in spatial and temporal variability within and/or among populations (Hess and Matala 2013; Hess *et al.* 2016a; Matala *et al.* 2017).

In the Columbia River Basin, Chinook Salmon have been studied extensively (e.g., Waples *et al.* 2004; Beacham *et al.* 2006; Narum *et al.* 2008b; Matala *et al.* 2011; Hecht *et al.* 2015), as have steelhead trout (Winans *et al.* 2004; Currens *et al.* 2009; Blankenship *et al.* 2011; Narum *et al.* 2011; Matala *et al.* 2016). The scope of Sockeye Salmon and Coho Salmon genetic monitoring has been comparatively limited but has received greater attention in recent years (Gustafson *et al.* 1997; Kozfkay *et al.* 2008; Iwamoto *et al.* 2012; Galbreath *et al.* 2014). Continued monitoring and evaluation of the genetic structure among salmon populations in the Columbia River Basin has guided managers in establishing and maintaining primary conservation units to protect fisheries resources. The delineation of such conservation units, including distinct population segment (DPS), evolutionarily significant unit (ESU), major population group (MPG), and viable salmonid population (VSP) is guided by a core set of criteria, including population ecology and viability, ancestry and descent, reproductive isolation, and genetic structure and local adaptation (Fraser and Bernatchez 2001; Fraser *et al.* 2011). Although an understanding of adaptive variation is critical to proper salmon management, the majority of genetic information available to managers is based on neutral genetic variation. Landscape genetics is an approach aimed at describing population differentiation relative to features in an organism's environment (Segelbacher 2010; Latch *et al.* 2011; Sepulveda-villet & Stepien 2012; Matala *et al.* 2014). Landscape genetics explores population differentiation relative to features in the environment such as migratory barriers (e.g., dams), or heterogeneous

habitats such as variation in local climates or temperatures (Dionne et al. 2008; Narum et al. 2008a; Micheletti et al. 2017). Although local adaptation may be inferred from landscape genetics (Olsen et al. 2010; Blankenship et al. 2011), inferences based primarily on neutral genetic differentiation risk incorrectly identifying the underlying processes affecting population distinctions (Funk et al. 2012; Landguth & Balkenhol 2012). Techniques such as outlier detection methods, and genome wide association studies (GWAS) based on DNA sequence variation provide evidence of non-neutral population structure or adaptive variation (markers associated with run timing; Hess et al. 2016b). Such applications in genetic monitoring allow a more resolved understanding of genetic differentiation beyond what can be concluded from neutral loci alone (Narum et al. 2010b; Matala et al. 2011; Ackerman et al. 2012, Bourret et al. 2013). Putative non-neutral population differentiation can then be interpreted in the context of contemporary risks and vulnerabilities (e.g., climate change) for salmonid populations in the Columbia River Basin, revealing highly correlative relationships between genetic variation and the physical environment (see Limborg et al 2011). This additional information may ultimately influence conservation criteria for delineating populations across diverse landscapes.

Project objectives, time line and harvest management questions:

Objective two of project #2008-907-00 (Genetic Assessment of Columbia River Stocks) describes efforts to evaluate genetic diversity among populations that will inform managers in the areas of harvest monitoring, and conservation monitoring. Our approach involves the collection, analysis, interpretation and distribution of genotypic data. These data are being compiled as species-specific reference baselines for characterizing Chinook Salmon, steelhead trout, and *O. nerka* population structure specific to the Columbia River Basin. Baselines were initially created from genotypes at single nucleotide polymorphism (SNP) loci, which are highly prolific in the genome and provide substantial coverage for linkage analyses (Moen et al. 2008). SNPs are amenable to superior high throughput capabilities and are relatively easily amplified and scored compared to other types of genetic markers, even with poor quality tissue (DNA) sources (Campbell and Narum 2008). Because SNPs are commonly found within or adjacent to coding and regulatory regions of a genome, corresponding allelic diversity and allele frequency variation are likely to be informative for understanding non-neutral influences (i.e. selection and local adaptation) on observed population structure. Large numbers of highly informative SNP loci have been discovered through our ongoing efforts using a next generation sequencing methods known as restriction-site associated DNA (RAD) sequencing (Miller et al. 2007; Baird et al. 2008; Hecht et al. 2013) and whole genome resequencing ([Whole Genome Resequencing: Poolseq Individual Barcode v1.0](#)). Our two primary objectives for utilizing SNP baselines to monitor salmon species in the Columbia River are 1) genetic stock identification (GSI) of natural-origin stocks, and 2) parentage based tagging (PBT), a large-scale, non-lethal tagging technology for monitoring and evaluating hatchery stocks. The collaborative, inter-agency application of GSI continues to provide invaluable monitoring capabilities to understand relative stock proportions in sport, commercial and tribal harvests, as well as monitoring of stock specific run-timing at Bonneville Dam, Lower Granite Dam and other fish weirs in the basin. Moreover, GSI is being used concordantly with PBT to monitor trends in hatchery production, harvest of hatchery fish, and population attributes of specific hatcheries (e.g., stray rates, survival/mortality, migratory behavior, hatchery/wild interactions). Additionally, our genetic baselines are being used to characterize populations in archival studies, to inform efforts to reintroduce fish into extirpated regions within historic ranges, and in domestication studies. In step with technological

advancements, further geographic coverage, and continuing marker development, our efforts continue to expand. Our most recent results will be reported on an annual basis, and data will be routinely uploaded to the FishGen.net database (<http://www.fishgen.net/home.aspx>) as a repository for data sharing and collaboration.

Methods

Baseline sampling and protocols:

Our previously established baselines, comprised of putatively neutral SNPs (e.g. 180 loci for *O. mykiss*), have been well characterized and have been used extensively for genetic stock identification (GSI) as described in Hess et al. (2015) and Hasselman et al. (2016). Our most recent efforts focus on expanding genetic characterizations throughout the basin that will provide information about adaptive potentials and natural selective forces contributing to stock structure. Next generation sequencing technologies continue to be employed by the CRITFC genetics lab in order to expand SNP panels for Chinook Salmon, Coho Salmon, Sockeye Salmon, steelhead trout and Pacific lamprey (Hess et al. 2016c). Methods for genotyping by sequencing (GT-seq) are described in Campbell et al. (2015), and protocols for whole genome resequencing are shown below. Detailed laboratory methods are available in Monitoring Methods:

<https://www.monitoringmethods.org/Protocol/Details/230> (ID#230; owner Matthew Campbell) and Hess et al. (2012). Methods for whole genome resequencing ([Whole Genome Resequencing: Poolseq Individual Barcode v1.0](#)) utilize PoolParty v0.8 (Micheletti and Narum 2018), a bioinformatic pipeline based on bash and R to identify and process SNPs. High-quality SNPs were then used to assess population structure based on principal component analyses (PCA) and pairwise genetic distance displayed in neighbor joining (NJ) trees using the ‘adegenet’ package in R (Jombart & Ahmed 2011) or in GenAlEx v6.5 (Peakall and Smouse 2006).

Expansion and status of reference baselines for GSI:

Our three primary goals for expanding, maintaining, and evaluating each species-specific baseline are threefold. First, we used genetic stock identification (GSI) analyses for monitoring of fishery returns through the migratory corridor of the Columbia River, including harvest GSI in the lower Columbia River, and fish passage GSI at Bonneville and Lower Granite dams (see sections 3 & 4 of this report). Second, PBT broodstock sampling and genotyping of Columbia River Basin hatcheries has been updated to include the most recent years for continued evaluation of hatchery stock composition in various fisheries and to more accurately account for abundance of natural-origin stocks by identifying unmarked hatchery fish. Third, the sequence/genotypic data are also being applied in various analyses to evaluate selection, including investigations of landscape genetics, and adaptive differentiation among populations.

The GT-seq primer pools developed from RAD-seq data are currently being used for all high throughput genotyping projects, including 5 target species: steelhead trout, Chinook Salmon, Coho Salmon, Sockeye Salmon, and Pacific Lamprey. Currently we do not utilize reference baselines for GSI of either Coho Salmon or Pacific Lamprey, and in 2019 there were no updates to the reference populations in our GSI baseline for Chinook Salmon and Sockeye Salmon. The most recent marker panels are described in Section 1. Testing of the reference baseline for GSI based on the expanded SNP marker panel for steelhead trout is described in Section 3.

In addition, we are using whole genome resequencing methods that pool samples (Pool-seq; Schlotterer et al. 2014) to generate reference baselines with millions of SNPs for several species, Chinook Salmon (Table 3), Coho Salmon (Table 4), and *O. mykiss* (Table 5). Allele frequencies from millions of SNPs will be analyzed to detect statistically significant regions of the genome associated with specific traits or adaptation to environmental factors. Putatively neutral regions of the genome are also useful for standard phylogeny and demographic analyses of populations. Therefore, there are multiple objectives that can be accomplished with these Pool-seq reference baselines including identifying novel SNPs that can be used for improved accuracy and precision of future GSI and PBT applications. These reference baselines comprised of millions of SNPs thus fit two of the main objectives for this project: SNP discovery and expand and create baselines.

Table 3. Creation of a Chinook Salmon SNP baseline by whole genome resequencing of pools of samples

Pool Seq Library ID	Number of reads (R1)	Mean Coverage	ID	Collection	(n)	Lineage	Reporting Groups
L-0730	438,161,235	35.02	OTS01	Big Creek Tule	71	Rogue	01_YOUNGS
L-0731	203,076,142	16.26	OTS03	Kalama R spring-run	93	LC	02_WCASSP
L-1037	203,899,979	16.07	OTS05	Elochoman R fall-run	86	LC	03_WCASFA
L-0732	241,863,213	14.64	OTS06	Lewis R fall-run	68	LC	03_WCASFA
L-1067	214,421,199	19.45	OTS12	White Salmon fall-run	78	LC	05_SPCRTU
L-0736/L-0736_2	751,365,978	64.28	OTS13	Spring Creek NFH tule fall-run	94	LC	05_SPCRTU
L-0737	248,031,203	21.27	OTS16	Warm Springs R spring-run	93	ST	07_DESCSP
L-0756	148,429,490	11.86	OTS17	John Day North Fork-Main Stem	52	ST	08_JOHNDR
L-0739	479,879,641	36.47	OTS18	Middle Fork John Day R spring-run	36	ST	08_JOHNDR
L-0740	139,194,175	9.99	OTS19	North Fork John Day R spring-run	39	ST	08_JOHNDR
L-0741_A	196,168,023	12.61	OTS20	American R spring-run	62	ST	09_YAKIMA
L-0906/L-0733/L-0733_T est/L-0876	208,041,237	16.89	-	Methow R spring-run	87	ST	10_UCOLSP
L-0742	340,007,427	14.54	OTS22	Winthrop NFH spring-run	83	ST	10_UCOLSP
L-0743	277,739,324	24.25	OTS24	Wenatchee R spring-run	51	ST	10_UCOLSP
L-0745	221,784,271	16.25	OTS26	Tucannon River spring-run	52	ST	11_TUCANO
L-1038	205,605,681	17.82	OTS28	Lostine R spring-run	41	ST	12_HELLSC
L-0747	279,162,633	21.60	OTS29	Grande Ronde R spring-run	30	ST	12_HELLSC

L-1069	195,959,654	16.30	OTS32	Red R spring-run	45	ST	12_HELLSC
L-0749	468,738,121	36.61	OTS33	Powell R spring-run	94	ST	12_HELLSC
L-0748	212,015,624	18.42	OTS30	Imnaha R spring-run	76	ST	12_HELLSC
L-1068	200,911,545	15.92	OTS31	Lolo Cr spring-run	48	ST	12_HELLSC
L-1087	194,515,174	17.94	OTS34	Rapid R Hatchery	96	ST	12_HELLSC
L-1070	217,481,702	19.89	OTS35	South Forth Salmon R spring-run	45	ST	13_SFSALM
L-0751	248,736,026	18.17	OTS37	Secesh R spring-run	90	ST	13_SFSALM
L-0752	570,363,043	49.47	OTS38	Chamberlain Cr spring-run	78	ST	14_CHMBLN
L-0753	238,267,170	10.02	OTS39	Big Cr spring-run	48	ST	15_MFSALM
L-1059	248,813,713	21.73	OTS41	Loon Cr spring-run	42	ST	15_MFSALM
L-0754	189,279,625	14.94	OTS43	Bear Valley Cr spring-run	33	ST	15_MFSALM
L-0755	218,687,602	20.14	OTS45	Marsh Cr spring-run	44	ST	15_MFSALM
L-1085	207,691,679	14.92	OTS40	Camas Cr spring-run	60	ST	15_MFSALM
L-1086	608,135,302	54.51	OTS42	Sulphur Cr spring-run	35	ST	15_MFSALM
L-1105	210,570,014	18.39	OTS44	Capehorn Cr spring-run	60	ST	15_MFSALM
L-0757	188,391,901	17.82	OTS47	Lemhi R spring-run	95	ST	16_UPSALM
L-0758	141,386,850	9.71	OTS48	Pahsimeroi R spring-run	92	ST	16_UPSALM
L-1057	203,885,206	18.56	OTS49	East Fork Salmon R spring-run	96	ST	16_UPSALM
L-1080	243,282,312	20.16	OTS50	Salmon R spring-run	61	ST	16_UPSALM
L-1060	239,722,545	14.02	OTS51	West Fork Yankee Fork spring-run	93	ST	16_UPSALM
L-0760	276,701,035	18.50	OTS53	Sawtooth Hatchery weir spring-run	91	ST	16_UPSALM
L-1106	208,349,688	20.87	OTS52	Valley Cr spring-run	42	ST	16_UPSALM
L-0761	503,337,440	32.75	OTS55	lower Yakima R fall-run	46	OT	18_UCOLSF
L-0762/L-0966/L-0967	476,295,447	32.01	OTS57	Wenatchee R summer-run	122	OT	18_UCOLSF
L-0763/L-0877/L-0897/L-0904	482,651,780	24.78	OTS59	Methow R summer-run	136	OT	18_UCOLSF
L-0764	215,478,389	17.03	OTS60	Lyons Ferry weir fall-run	92	OT	19_SRFALL

Table 4. Creation of a Coho Salmon SNP baseline by whole genome resequencing of pools of samples

Stock	n	Location	Region (BPA subbasin)	Lineage
Wenatchee	282	Leavenworth NFH	Wenatchee	Oki-Lower Columbia
Bonneville Dam	175	Bonneville Dam	Mixed	Mixed

Table 5. Creation of *O. mykiss* SNP baseline by whole genome resequencing of pools of samples

Lineage	Reporting Group	Major Subbasin	Subbasin	Reach	Sites	N	Mapped Reads	Prop. Geno. @ Dep.*	
Coastal	02_LOWCOL	L. Columbia	Cowlitz	Cowlitz	Cowlitz R, Coweeman R	95	257,455,270	0.61	¹
Coastal	02_LOWCOL	L. Columbia	Hood	EF Hood	EF Hood R	48	260,113,702	0.62	¹
Coastal	02_LOWCOL	L. Columbia	Hood	EF Hood	Parkdale Fish Facility	83	219,546,080	0.51	¹
Coastal	02_LOWCOL	L. Columbia	Hood	WF Hood	WF Hood R	79	212,615,249	0.45	²
Coastal	02_LOWCOL	L. Columbia	Lewis	Lewis	EF Lewis R	78	325,463,369	0.65	¹
Coastal	03_SKAMAN	L. Columbia	Washougal	Washougal	Skamania Stock	60	250,089,125	0.58	¹
Coastal	04_WILLAM	L. Columbia	Willamette	Willamette	Eagle Cr	61	254,529,741	0.58	¹
Coastal	04_WILLAM	L. Columbia	Willamette	Willamette	Little Rock, Mad Crks	50	265,050,782	0.55	¹
Coastal	05_BWSALM	L. Columbia	Big White Salmon	Big White Salmon	Big White Salmon R	95	194,626,391	0.48	²
Inter-mediate	06_KLICKR	L. Columbia	Klickitat	Klickitat	Swale, Brush, Dead Canyon, Synder, White, Tepee, Fish Lake Crks	95	227,020,456	0.52	²
Inland	07_MGILCS	L. Columbia	Columbia Gorge	Columbia Gorge	Mill Cr	96	232,750,154	0.56	¹
Inland	07_MGILCS	L. Columbia	Columbia Gorge	Columbia Gorge	Fifteenmile Cr	92	170,322,546	0.62	³
Inland	07_MGILCS	L. Columbia	Deschutes	Deschutes	Warm Springs R	95	262,227,329	0.61	¹
Inland	07_MGILCS	L. Columbia	John Day	John Day	MF John Day R, Granite, Rudio, Wall, Trail Crks	95	285,409,446	0.58	¹
Inland	07_MGILCS	L. Columbia	John Day	MF John Day	Upper John Day R, Beech, Belshaw, Canyon Crks	69	246,704,439	0.57	¹
Inland	07_MGILCS	L. Columbia	John Day	SF John Day	Murderers, Deer, Black Canyon Crks	96	224,529,764	0.57	¹
Inland	07_MGILCS	L. Columbia	Middle Columbia	Middle Columbia	Rock, Squaw Crks	91	216,427,544	0.61	³
Inland	07_MGILCS	L. Columbia	Umatilla	Umatilla	Umatilla R	70	236,157,293	0.54	²
Inland	07_MGILCS	L. Columbia	Walla Walla	Walla Walla	Walla Walla R, Touchet R, Yellowhawk Cr	95	158,223,069	0.48	³
Inland	07_MGILCS	Snake	Asotin	Asotin	Asotin Cr	60	227,071,819	0.57	¹
Inland	07_MGILCS	Snake	Asotin	Asotin	George Creek	58	321,868,839	0.66	¹
Inland	07_MGILCS	Snake	Clearwater	L Clearwater	Little Bear Cr	46	267,109,802	0.60	¹
Inland	07_MGILCS	Snake	Clearwater	L Clearwater	Sweetwater Cr	51	224,853,521	0.53	¹
Inland	07_MGILCS	Snake	Clearwater	L Clearwater	Potlatch R	50	217,298,385	0.51	¹
Inland	07_MGILCS	Snake	Grande Ronde	Grande Ronde	Big Canyon Cr	95	251,726,901	0.59	¹

Inland	07_MGILCS	Snake	Grande Ronde	Grande Ronde	Catherine Cr	91	199,865,574	0.53	²
Inland	07_MGILCS	Snake	Grande Ronde	Grande Ronde	Joseph Cr	88	240,989,824	0.52	¹
Inland	07_MGILCS	Snake	Grande Ronde	Grande Ronde	upper Grande Ronde R	58	190,569,290	0.61	³
Inland	07_MGILCS	Snake	Imnaha	Imnaha	Gumboot Cr	38	233,911,089	0.54	¹
Inland	07_MGILCS	Snake	Imnaha	Imnaha	Lightning Cr	95	319,903,348	0.66	¹
Inland	07_MGILCS	Snake	Imnaha	Imnaha	Little Sheep Cr	76	235,566,015	0.53	¹
Inland	07_MGILCS	Snake	Lower Snake	Lower Snake	Alpowa Cr	53	287,842,645	0.63	¹
Inland	07_MGILCS	Snake	Salmon	L Salmon	Little Salmon R, Hazard, Boulder Crks	95	248,041,850	0.59	¹
Inland	07_MGILCS	Snake	Salmon	L Salmon	White Bird Cr	50	200,302,136	0.46	²
Inland	07_MGILCS	Snake	Tucannon	Tucannon	Tucannon R	42	214,476,660	0.55	¹
Inland	07_MGILCS	Snake	Umatilla	Umatilla	Minthorn Springs	74	222,263,607	0.54	¹
Inland	08_YAKIMA	U. Columbia	Yakima	Yakima	Naches R, Nile Cr	38	181,114,635	0.49	³
Inland	08_YAKIMA	U. Columbia	Yakima	Yakima	Satus R	67	208,951,362	0.49	²
Inland	08_YAKIMA	U. Columbia	Yakima	Yakima	Teaway R, Big Cr, Roza Dam	95	384,909,116	0.69	¹
Inland	09_UPPCOL	U. Columbia	Entiat	Entiat	Entiat R	43	336,470,474	0.65	¹
Inland	09_UPPCOL	U. Columbia	Methow	Methow	Winthrop NFH	93	180,992,366	0.46	²
Inland	09_UPPCOL	U. Columbia	Okanogan	Okanogan	Bonaparte, Salmon, Omak Crks	95	278,354,930	0.58	¹
Inland	09_UPPCOL	U. Columbia	Wenatchee	Wenatchee	Chiwaukum Cr	54	227,418,107	0.56	¹
Inland	10_SFCLWR	Snake	Clearwater	SF Clearwater	Clear Cr	61	243,826,940	0.59	¹
Inland	10_SFCLWR	Snake	Clearwater	SF Clearwater	Crooked R	86	216,426,527	0.51	¹
Inland	10_SFCLWR	Snake	Clearwater	SF Clearwater	Tenmile Cr	60	288,038,275	0.64	¹
Inland	11_UPCLWR	Snake	Clearwater	Lochsa	Canyon, Deadman Crks	68	285,197,311	0.61	¹
Inland	11_UPCLWR	Snake	Clearwater	Lochsa	upper Lochsa R	72	129,797,910	0.57	³
Inland	11_UPCLWR	Snake	Clearwater	Selway	Little Clearwater R	65	236,963,840	0.55	¹
Inland	11_UPCLWR	Snake	Clearwater	Selway	upper Selway R	51	111,568,814	0.49	³
Inland	11_UPCLWR	Snake	Clearwater	Selway	White Cap Cr	72	264,031,261	0.60	¹
Inland	12_SFSALM	Snake	Salmon	SF Salmon	Johnson Cr	95	201,270,498	0.48	²
Inland	12_SFSALM	Snake	Salmon	SF Salmon	Lick Cr	70	200,066,360	0.56	²
Inland	12_SFSALM	Snake	Salmon	SF Salmon	Secesh R	30	263,042,376	0.60	¹
Inland	12_SFSALM	Snake	Salmon	SF Salmon	East Fork SF Salmon R	53	126,942,745	0.56	³
Inland	13_MFSALM	Snake	Salmon	MF Salmon	Bargamin Cr	60	189,953,336	0.53	²
Inland	13_MFSALM	Snake	Salmon	MF Salmon	Big, Rush Crks	95	328,230,174	0.67	¹
Inland	13_MFSALM	Snake	Salmon	MF Salmon	Loon Cr	51	320,069,394	0.65	¹
Inland	13_MFSALM	Snake	Salmon	MF Salmon	Marsh Cr	60	334,274,383	0.67	²
Inland	14_UPSALM	Snake	Salmon	NF Salmon	Sawtooth Hatchery	47	241,483,030	0.57	¹
Inland	14_UPSALM	Snake	Salmon	NF Salmon	EF Salmon R	51	304,068,477	0.64	¹
Inland	14_UPSALM	Snake	Salmon	NF Salmon	Yankee Fork Salmon R	58	191,268,742	0.52	²

Inland	14_UPSALM	Snake	Salmon	NF Salmon	Morgan Cr	39	109,730,025	0.50	³
Inland	14_UPSALM	Snake	Salmon	NFSalmon	Lemhi R, Bear Valley Cr	95	225,800,707	0.51	²

*proportion of draft genome covered at a given depth of 10X:

¹ 100bp trim, min depth 15; ² 50bp trim, min depth 15; ³ 50bp trim, min depth 10

Results

Parentage Based Tagging (PBT) update

PBT began with Chinook Salmon and steelhead hatchery stocks in the Snake River basin of Idaho (2008-present; Steele et al., 2013; Steele et al., 2015). However, we have expanded PBT coverage (Figure 10, Figure 14, Appendix 6, Appendix 7, and Appendix 8) to include Chinook Salmon, steelhead, and Coho Salmon broodstocks in all hatcheries above Bonneville Dam using expanded SNP panels of 343 loci for Chinook Salmon, 368 loci for steelhead trout and 257 loci for Coho Salmon. Each year the expansion effort is integrated with existing PBT baselines as data comes available (e.g., Chinook salmon, Appendix 4).

Adopting PBT to the broader Col. River basin facilitates our ability to genetically track millions of salmonids and provide opportunities to address a variety of parentage-based research and management questions, including stock contributions to fisheries (Byrne et al., 2015), estimates of stock-specific abundance and run-timing at dams (Hess et al., 2016c; Vu et al. 2015), and use of thermal refugia during migration (Hess et al., 2016a).

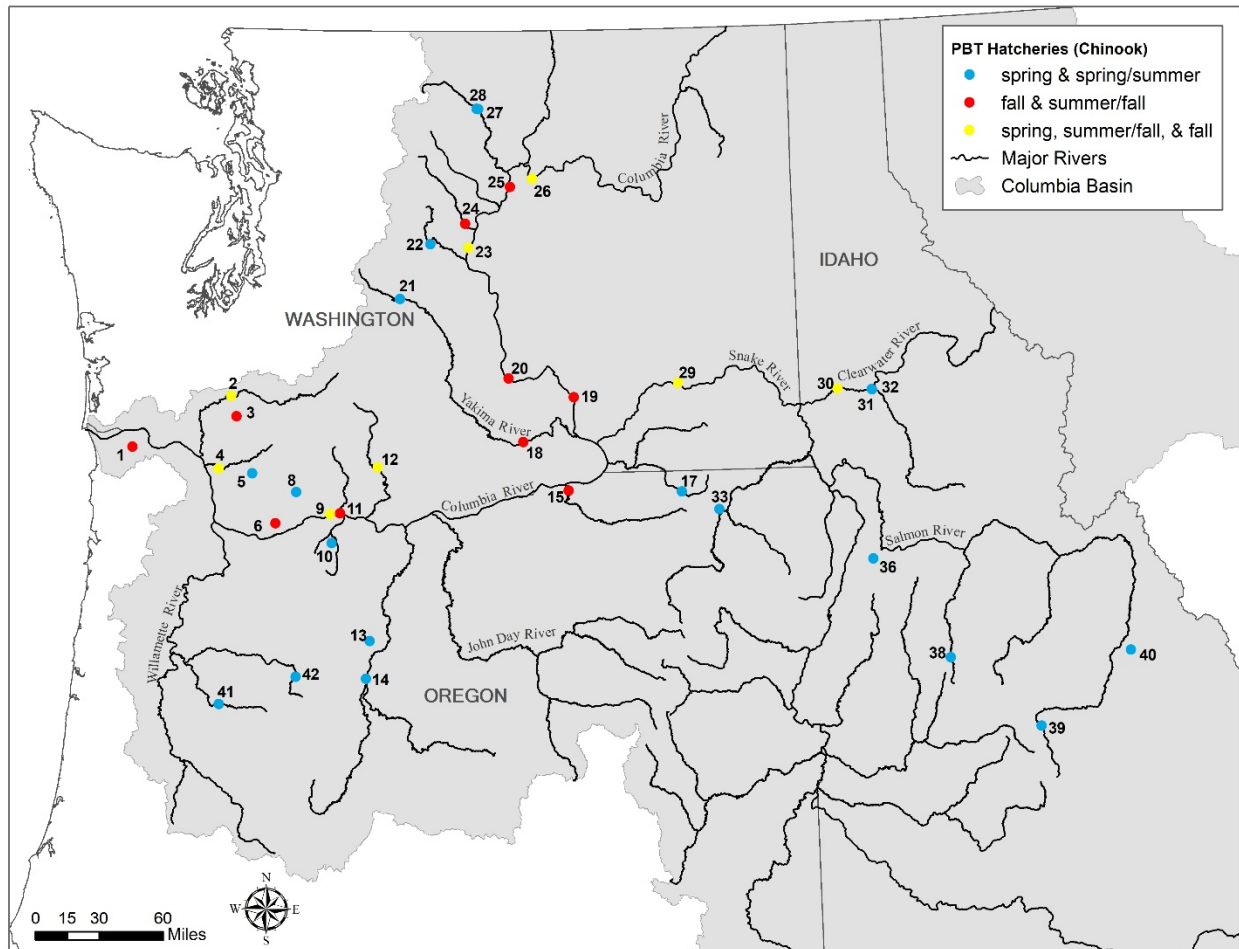


Figure 10. Chinook Salmon, PBT hatcheries. Numbers correspond to map ID and hatchery descriptions (Appendix 6).

The PBT tagging technology has been implemented through annual hatchery broodstock sampling to create a temporally structured parental genotype baseline. As adult fish return to participating hatcheries in the Columbia River basin, broodstock are sampled by collection of fin tissue during hatchery spawning. Required data for PBT sampling includes a hatchery record of phenotypic sex and spawn date. Additional and optional information was collected at some hatcheries when resources allowed, including fork length, and mated cross records of male and female broodstock individuals. The PBT baseline expansion during the 2019 report period included spawn year 2017 for some lower Columbia River hatcheries, but primarily broodyear 2018 was genotyped for most hatcheries. The total numbers of fish genotyped for PBT baselines in 2019 included $n=16,529$ spring Chinook Salmon, $n=2,563$ upper Columbia summer Chinook Salmon, $n=28,514$ fall Chinook Salmon (Appendix 4), $n=4,722$ steelhead trout (Appendix 5), and $n=1,758$ Coho Salmon (Appendix 5). DNA was extracted using modified Chelex extractions and Qiagen DNeasy 96 kits. Extracted genomic DNA was genotyped at 343 SNP loci for Chinook Salmon, 368 SNP loci for steelhead trout, and 257 SNP loci for Coho Salmon using a GTseq protocol: (<https://www.monitoringresources.org/Document/Method/Details/5446>).

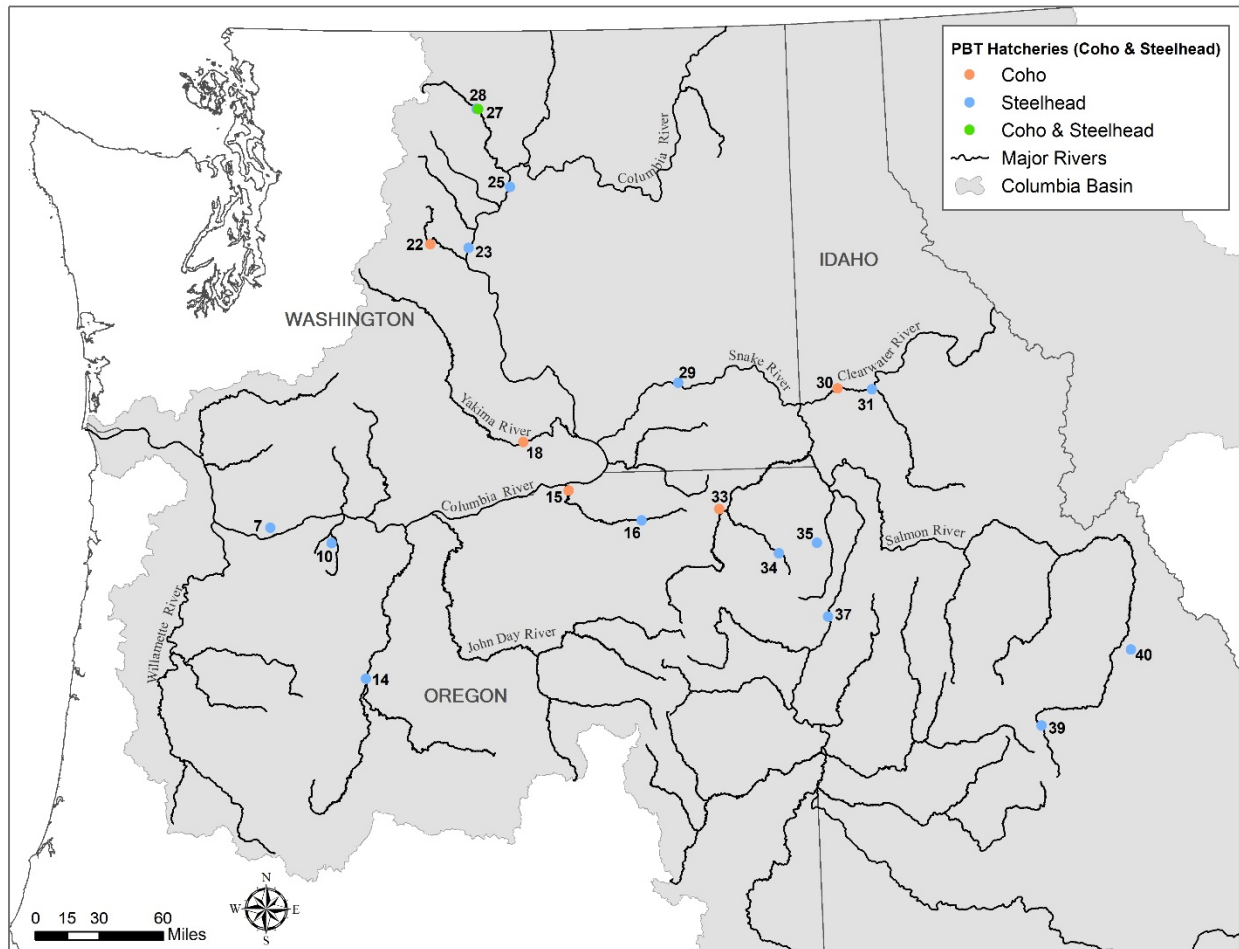


Figure 11. Steelhead and Coho Salmon, PBT hatcheries. Numbers correspond to map ID and hatchery descriptions (Appendix 7, Appendix 8).

Hatchery offspring that are subsequently sampled either as juveniles or adults (e.g., in a fishery) are then PBT assigned back to spawned parents which provides the individual age and specific hatchery of origin for each offspring.

Discussion

Over the course of the Genetic Assessment project we have compiled extensive data sets of SNP genotypes for Chinook Salmon steelhead trout and *O. nerka*, and continue to improve resources for Coho Salmon and Pacific Lamprey. The populations we have evaluated span diverse regions in the Columbia River Basin (including the Snake River Basin). We continue to update and maintain SNP reference baselines for evaluation of these species in future generations. This long-term strategy informs harvest management and assures the greatest likelihood of discerning temporal variation among reproductively distinct species aggregates (Waples 1991), while monitoring population viability related to demographic trends that occur locally and/or regionally. Philopatry (Quinn et al. 1991, Hendry et al. 2003) and hatchery supplementation activities (Ford et al. 2006; Hard & Heard 1999) play a major role in how genetic divergence and differentiation is distributed geographically. For managing sustainable fisheries, it is necessary to understand the magnitude of influence that these and other factors have on our ability to differentiate populations, both qualitatively (phenotypes; landscapes) and quantitatively (e.g., genetic stock identification). This becomes particularly important where mixed stock fisheries may consist of both ESA listed and unlisted populations, and where differential harvest may have the greatest impact on specific populations. Our current efforts have largely focused on expanding numbers of markers and non-neutral markers associated with maturation. However, loci for adaptive divergence (landscape genetics) remains as one of our primary objectives for strengthening our understanding of non-neutral genetic variation among populations. Data collected through whole genome resequencing and GT-seq techniques has yielded large numbers of potential SNPs, and demonstrated their utility for characterizing adaptive variation, and identifying environmental and life history related variables that are likely to have significant influence on allele frequencies (e.g., precipitation, temperature, run-timing.; Hecht et al. 2015, Hess et al. 2016b; Micheletti et al. 2017; Narum et al. 2018; Micheletti et al. 2018). The expansion efforts reported here also provided improved ability to differentiate stocks on regional and local scales through application of GSI and PBT methods.

Collections of *O. tshawytscha*, *O. mykiss*, and *O. nerka* have been chosen for baseline expansion based on availability, novelty, and in accordance with our goal of reaching complete coverage of extant stocks within the Columbia River Basin. Priority collections for all three species have been identified as those important to basin-wide harvest and hatchery management, particularly in tribal fisheries. This includes major supplementation stocks for all three species: lower Columbia, ocean-type, and stream-type lineages of Chinook Salmon, inland and coastal lineages of steelhead trout, and the anadromous (Sockeye Salmon) and land-locked (kokanee) forms of *O. nerka*. Species-specific reference baselines may include life history variants such as potentially distinct populations of resident *O. mykiss* (Narum et al. 2008a; Narum et al. 2011). The application of GSI in fisheries continues to inform managers on several fronts, including: harvest management, abundance estimates, life history distinctions and conservation needs. Moreover, PBT is being used for multiple purposes including validation of assigned origins using GSI. In fact, PBT frequently reveals substantial numbers of unmarked hatchery-origin fish that are incorrectly identified as wild in the field. Future efforts for baseline expansion include compiling allele frequencies for millions of SNPs from whole genome resequencing that can be drawn from at any time should the need for more markers be necessary. An example of such need is basin-wide coverage to account for stock transfers or reintroductions throughout the basin.

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pop	subbasin	location	latitude	longitude
<u>Coastal Lineage</u>				
Grays	Grays	Gray R.	46.2121	-123.3634
Elochoman	Elochoman	Elochoman R.	46.2267	-123.3304
Mill	Elochoman	Mill Cr.	46.1885	-123.1773
Abernathy	Elochoman	AFTC weir	46.2256	-123.1481
Germany	Elochoman	Germany Cr.	46.1901	-123.1238
Cowlitz	Cowlitz	Cowlitz R. at Barrier Dam	46.5026	-122.5881
Kalama (SUM)	Kalama	Kalama R.	46.0329	-122.8703
Kalama (WIN)	Kalama	Kalama R.	46.0329	-122.8703
Lewis	Lewis	North Fork Lewis @ Merwin Dam	45.9568	-122.5551
Eagle	Willamette	Clackamas R. - Eagle Cr.	45.3514	-122.3840
Clackamas (SUM)	Willamette	Clackamas R. - North Fork Dam	45.2417	-122.2817
Clackamas (WIN)	Willamette	Clackamas R. - Eagle Cr.	45.3514	-122.3840
	Willamette	Clackamas R. - Timber Park	45.2994	-122.3506
	Willamette	Columbia R. - mainstem	45.3724	-122.6044
N. Santiam	Willamette	Rock Cr.	44.7467	-122.3953
S. Santiam	Willamette	Wiley Cr.	44.4150	-122.6739
Sandy	Sandy	Still Cr.	45.3309	-121.9158
Washougal	Washougal	Washougal R.	45.3909	-122.0960
BWS Rattlesnake	Big White Salmon	Rattlesnake Cr.	45.7993	-121.4843
	Big White Salmon	BWS screw trap	45.7476	-121.5211
BWS mainstem	Big White Salmon	mainstem (Dam6 relocation)	45.7993	-121.4843
	Big White Salmon	BWS screw trap	45.7476	-121.5211
Skamania	Washougal	Skamania Hatchery	45.6227	-122.2175
<u>Inland Lineage</u>				
Fifteenmile	Fifteenmile	Fifteenmile Cr.	45.4509	-121.1244
Deschutes	Deschutes	Deschutes R. - Trout Cr.	44.8214	-121.0872
	Deschutes	Warm Springs R.	44.8609	-121.2444
	Deschutes	Warm Springs R. - Mill Cr.	44.8647	-121.4222
	Deschutes	Warm Springs R. - Mill Cr.	44.8647	-121.4222
Klickitat	Klickitat	Dillacort Cr.	45.7414	-121.2222
	Klickitat	Wheeler Cr.	45.7951	-121.1928
	Klickitat	Swale Cr.	45.8091	-121.0652
	Klickitat	Snyder Cr.	45.8247	-121.1565
	Klickitat	Little Klickitat R.	45.8271	-120.8181
	Klickitat	Bowman Cr.	45.8452	-121.0421
	Klickitat	Dead Canyon	45.9423	-121.1439
	Klickitat	Summit Cr.	45.9876	-121.1255
	Klickitat	White Cr.	46.0133	-121.1500
	Klickitat	Trout Cr.	46.0378	-121.1994
JohnDay	John Day	Beech Cr.	44.4113	-119.1160
	John Day	Black Canyon Cr.	44.3314	-119.5657
	John Day	Murderer's Cr.	44.3173	-119.5310
	John Day	Deer Cr.	44.1890	-119.5139
	John Day	Middle Fork John Day	44.9129	-119.2964
	John Day	Fox Cr.	44.6156	-119.2941
	John Day	Belshaw Cr.	44.4370	-119.2930
	John Day	Desolation Cr.	44.9397	-118.8379
	John Day	Camp Cr.	44.6891	-118.7966
	John Day	Clear Cr.	44.5888	-118.5068
	John Day	Granite Cr.	44.8383	-118.4770
	John Day	Granite Cr.	44.8383	-118.4770
Umatilla	Umatilla	Iskuulpa Cr.	45.6994	-118.3965
Ahtanum	Yakima	Ahtanum R.	46.5318	-120.6815
Cowiche	Yakima	Cowiche Cr.	46.6288	-120.5708
Naches	Yakima	Little Rattlesnake Cr.	46.7680	-121.0464
	Yakima	N.F. Little Naches R.	47.1090	-121.3201

	Yakima	Pile Up Cr.	46.3274	-121.1537
	Yakima	Nile Cr.	46.8618	-121.0489
	Yakima	Quartz Cr.	45.9457	-120.5097
Satus	Yakima	Dry Cr.	46.1929	-120.6138
	Yakima	Satus Cr.	46.0432	-120.5716
Toppenish	Yakima	Toppenish Cr.	46.3227	-120.8749
Wenatchee	Wenatchee	Levenworth NFH	47.5591	-120.6742
	Wenatchee	Nason Cr.	47.8019	-120.7150
	Wenatchee	Wells Hatchery (LNFH Stock)	47.5591	-120.6742
Methow	Methow	Methow R.	48.0495	-119.9012
Entiat	Entiat	Entiat R. - screw trap	47.6641	-120.2417
Omak	Okanogan	Omak Cr. Adult Weir	48.3957	-119.5043
Touchet	Walla Walla	Touchet R.	46.0340	-118.6836
Dworshak	Clearwater	Dworshak National Fish Hatchery	46.5025	-116.3205
Lochsa	Clearwater	Fish Cr.	46.3316	-115.3461
S. F. Clearwater	Clearwater	South Fork Clearwater R.	46.5061	-116.5491
Catherine	Grande Ronde	Catherine Cr.	45.3070	-117.8660
Joseph	Grande Ronde	Joseph Cr.	46.0416	-117.0017
UpperGR	Grande Ronde	Upper Grande Ronde Adult Trap	45.7314	-117.8640
Wenaha	Grande Ronde	Wenaha R.	45.9453	-117.4513
Lightning	Imnaha	Lightning Cr. Weir	45.6554	-116.7265
Rapid	Salmon	Little Salmon R. - Rapid R.	45.3547	-116.3915
N. F. Salmon	Salmon	North Fork Salmon R.	45.4099	-113.9919
Pahsimeroi	Salmon	Pahsimeroi Fish Hatchery	44.6845	-114.0404

total

305

306

Note: Each locality is identified by the lineage, subbasin- specific location (e.g. tributary) with geographic coordinates, origin, and sample size (n).

Map ID	Spawning hatchery	Run type	Lineage	Region	Latitude	Longitude	2019 genotyping		
							Year	Sampled	Completed
	Klaskanine (NF brood)	fall (tule)	LC	Col.	46.09	-123.718	na	na	na
	Klaskanine (SF brood)	fall (tule)	LC	Col.	46.09	-123.718	na	na	na
1	Big Creek	fall (tule)	LC	Col.	46.147	-123.581	2015	2018	1886
2	Cowlitz Salmon	fall (tule)	LC	Col.	46.511	-122.629	2015	2017	1929
2	Cowlitz Salmon	spring	LC	Col.	46.511	-122.629	2015	2017	1649
3	Toutle	fall (tule)	LC	Col.	46.375	-122.572	2015	2017	167
4	Kalama Falls	fall (tule)	LC	Col.	46.017	-122.733	2016	2017	3460
4	Kalama Falls	spring	LC	Col.	46.017	-122.733	2015	2017	342
5	Lewis River	spring	LC	Col.	45.937	-122.616	2015	2017	964
	Clackamas	spring	LC	Col.	45.296	-122.362	na	na	na
42	Marion Forks	spring	LC	Col.	44.612	-121.948	2018	2018	920
41	South Santiam	spring	LC	Col.	44.416	-122.675	2015	2018	3159
	McKenzie	spring	LC	Col.	44.118	-122.637	na	na	na
	Willamette	spring	LC	Col.	43.745	-122.444	na	na	na
6	Washougal	fall (tule)	LC	Col.	45.653	-122.169	2015	2017	1177
	Bonneville, Tanner Cr.	fall (tule)	LC	Col.	45.633	-121.957	na	na	na
11	Spring Creek NFH	fall (tule)	LC	Col.	45.728	-121.544	2015	2018	6524
9	Little White Salmon NFH	fall	IOT	Col.	45.719	-121.645	2013	2018	1743
15	Umatilla	fall	IOT	Col.	45.913	-119.552	2012	2018	968
12	Klickitat	fall	IOT	Col.	46.041	-121.183	2018	2018	491
18	Prosser	fall	IOT	Col.	46.215	-119.76	2012	2018	87
19	Ringold Springs	fall	IOT	Col.	46.514	-119.259	2016	2018	455
20	Priest Rapids	fall	IOT	Col.	46.647	-119.899	2012	2018	6937
29	Lyons Ferry	fall	IOT	Snake	46.598	-118.226	2011	2018	1879
30	Nez Perce Tribal	fall	IOT	Snake	46.52	-116.66	2011	2018	811
23	Eastbank	summer	IOT	Col.	47.53	-120.293	2012	2018	723
24	Entiat NFH	summer	IOT	Col.	47.698	-120.323	2013	2018	309
25	Wells	summer	IOT	Col.	47.947	-119.871	2012	2018	777
26	Chief Joseph	summer	IOT	Col.	48.001	-119.647	2013	2018	754
9	Little White Salmon NFH	spring	IST	Col.	45.719	-121.645	2013	2018	1644
15	Umatilla	spring	IST	Col.	45.913	-119.552	2012	2018	459
29	Lyons Ferry	spring	IST	Snake	46.598	-118.226	2008	2018	IDFG
30	Nez Perce Tribal	spring	IST	Snake	46.52	-116.66	2008	2018	188
23	Eastbank	spring	IST	Col.	47.53	-120.293	2012	2018	272
26	Chief Joseph	spring	IST	Col.	48.001	-119.647	2014	2018	229
8	Carson NFH	spring	IST	Col.	45.868	-121.974	2012	2018	850
10	Parkdale	spring	IST	Col.	45.525	-121.622	2012	2018	194
12	Klickitat	spring	IST	Col.	46.041	-121.183	2008	2018	197
13	Warm Springs NFH	spring	IST	Col.	44.861	-121.246	2012	2018	307
14	Round Butte	spring	IST	Col.	44.605	-121.277	2012	2018	696
21	Cle Elum SRF	spring	IST	Col.	47.187	-120.976	2012	2018	511
22	Leavenworth NFH	spring	IST	Col.	47.558	-120.674	2013	2018	721
27	Methow	spring	IST	Col.	48.477	-120.205	2012	2018	129
28	Winthrop NFH	spring	IST	Col.	47.558	-120.675	2013	2018	387
33	Lookingglass	spring	IST	Snake	45.732	-117.865	2008	2018	761
31	Dworshak NFH	spring	IST	Snake	46.504	-116.328	2008	2018	IDFG
32	Clearwater	spring	IST	Snake	46.504	-116.328	2008	2018	IDFG
36	Rapid River	spring/summer	IST	Snake	45.354	-116.394	2008	2018	1855
38	SF Salmon, McCall	spring/summer	IST	Snake	44.908	-116.117	2008	2018	IDFG
38	SF Salmon, Johnson Creek	spring/summer	IST	Snake	44.899	-115.492	2008	2018	95
40	Pahsimeroi	spring/summer	IST	Snake	44.684	-114.039	2008	2018	IDFG
39	Sawtooth	spring/summer	IST	Snake	44.15	-114.883	2008	2018	IDFG

* In 2017 adult broodstock for Umatilla were collected/spawned at Ringold Springs.
Note: The Map ID indicates the sites of the hatcheries corresponding to Figure 13. Genetic lineage is lower Col. (LC), interior ocean-type (IOT), and interior stream-type (IST). Year refers to the first year of PBT sampling for

313 each hatchery: na – not currently a PBT hatchery. For some hatchery PBT samples, genotyping efforts have begun
314 to backdate collections that were initially archived- the “sampled” field indicates which collection years were
315 genotyped in 2018. The project collaborators at Idaho Department of Fish and Game (IDFG) were responsible for
316 genotyping of Snake River hatcheries (see “completed” column).

317 **Appendix 5. Steelhead and Coho Salmon hatchery broodstock sampled for PBT baselines.**

								<u>2019 Genotyping</u>	
Map ID	Spawning hatchery	Run type	Lineage	Region	Latitude	Longitude	Year	Sampled	Completed
1	Big Creek	winter	coastal	Col.	46.147	-123.581	na	na	na
	Abernathy FTC	winter	coastal	Col.	46.226	-123.153	2012	archived	0
	Cowlitz Trout	winter	coastal	Col.	46.511	-122.629	na	na	na
4	Kalama Falls	winter	coastal	Col.	46.017	-122.733	na	na	na
	Merwin	winter	coastal	Col.	45.954	-122.564	na	na	na
	Clackamas	winter	coastal	Col.	45.296	-122.362	na	na	na
	Eagle Creek NFH	winter	coastal	Col.	45.276	-122.202	na	na	na
	Sandy	winter	coastal	Col.	45.407	-122.254	na	na	na
7	Skamania	summer/winter	coastal	Col.	45.652	-122.168	2013	2018	346
10	Parkdale	winter	coastal	Col.	45.525	-121.622	2012	2018	41
14	Round Butte	summer	inland	Col.	44.605	-121.277	2013	2018	906
15	Umatilla	summer	inland	Col.	45.913	-119.552	2012	2018	83
23	Eastbank	summer	inland	Col.	47.53	-120.293	2012	2018	145
25	Wells	summer	inland	Col.	47.947	-119.871	2013	2018	238
27	Methow (Twisp)	summer	inland	Col.	48.477	-120.205	2013	2018	na*
28	Winthrop NFH	summer	inland	Col.	48.477	-120.205	2012	2018	132
29	Lyons Ferry	summer	inland	Snake	46.598	-118.226	2009	2018	IDFG
34	Wallowa	summer	inland	Snake	45.418	-117.302	2009	2018	IDFG
37	Oxbow	summer	inland	Snake	44.971	-116.853	2008	2018	IDFG
31	Dworshak NFH	summer	inland	Snake	46.504	-116.328	2008	2018	1792
40	Pahsimeroi	summer	inland	Snake	44.684	-114.039	2008	2018	1039
39	Sawtooth	summer	inland	Snake	44.15	-114.883	2008	2018	IDFG
35	Little Sheep Creek	summer	inland	Snake	45.477	-116.928	2008	2018	IDFG

Coho

15	Umatilla	na	na	Col.	45.913	-119.552	2012	2018	589
18	Prosser	na	na	Col.	46.215	-119.76	2016	2018	579
22	Leavenworth NFH	na	na	Col.	47.558	-120.674	2012	2018	na
28	Winthrop NFH	na	na	Col.	48.477	-120.205	2012	2018	na
31	Dworshak NFH	na	na	Snake	46.504	-116.328	2012	2018	590

* Steelhead Methow Hatchery Twisp stock spawned at Winthrop NFH starting in 2017; not distinguished from Winthrop stock.

Note: The map ID indicates site locations corresponding with Figure 13. Genetic lineage is coastal or inland. Year refers to the first year of PBT sampling for each hatchery: na – not currently a PBT hatchery. Some 2017 PBT samples have been archived awaiting shipment to the Hagerman Genetics Laboratory as of the drafting of this report. The project collaborators at Idaho Department of Fish and Game (IDFG) were responsible for genotyping of Snake River hatcheries (see “completed” column). All Coho broodstocks sampled for PBT broodstock were designated for release of fish upstream of Bonneville Dam.

Appendix 6. Checklist of PBT broodstock collections that comprise the PBT baselines for Chinook salmon from 2008 through 2018.

Map num.	Hatchery	Species	Code	Run type	Lineage	Year											
						2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	
32	Clearwater Fish Hatchery	Chinook	OtsCLWH	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
32	Clearwater Fish Hatchery - Powell Facility	Chinook	OtsPOWP	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
31	Dworshak National Fish Hatchery	Chinook	OtsDWOR	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
33	Lookingglass Fish Hatchery - Catherine Creek	Chinook	OtsCTHW	Spring/Summer	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
33	Lookingglass Fish Hatchery - Grande Ronde	Chinook	OtsGRUW	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
33	Lookingglass Fish Hatchery - Imnaha River	Chinook	OtsIMNW	Spring/Summer	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
33	Lookingglass Fish Hatchery - Lookingglass Creek	Chinook	OtsLOOK	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
33	Lookingglass Fish Hatchery - Lostine River	Chinook	OtsLSTW	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery	Chinook	OtsLYON	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery - Tucannon River	Chinook	OtsTUCW a	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery	Chinook	OtsLYON_1	Fall	Interior ocean type	*	*	*	X	X	X	X	X	X	X	X	
38	McCall Fish Hatchery - Johnson Creek	Chinook	OtsJHNW	Spring/Summer	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
38	McCall Fish Hatchery - South Fork Salmon	Chinook	OtsMCCA	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
30	Nez Perce Tribal Fish Hatchery	Chinook	OtsNPFH_1	Fall	Interior ocean type	*	*	*	X	X	X	X	X	X	X	X	
30	Nez Perce Tribal Fish Hatchery	Chinook	OtsNPFH	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
40	Pahsimeroi Fish Hatchery	Chinook	OtsPAHH	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
36	Rapid River Fish Hatchery	Chinook	OtsRAPH	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
39	Sawtooth Fish Hatchery	Chinook	OtsSAWT	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
1	Big Creek Hatchery	Chinook	OtsBIG_fa	Fall	Lower Columbia	*	*	*	*	*	*	*	X	X	X	X	
8	Carson National Fish Hatchery	Chinook	OtsCAR_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
26	Chief Joseph Hatchery	Chinook	OtsCJH_sp	Spring	Interior stream type	*	*	*	*	*	*	X	X	X	X	X	
26	Chief Joseph Hatchery - Integrated	Chinook	OtsCJHint_su	Summer	Interior ocean type	*	*	*	*	*	X	X	X	X	X	X	
26	Chief Joseph Hatchery - Segregated	Chinook	OtsCJHseg_su	Summer	Interior ocean type	*	*	*	*	*	X	X	X	X	X	X	
2	Cowlitz Salmon	Chinook	OtsCOW_sp	Spring	Interior stream type	*	*	*	*	*	*	*	X	X	X	**	
2	Cowlitz Salmon	Chinook	OtsCOW_fa	Fall	Interior stream type	*	*	*	*	*	*	*	X	X	X	**	
23	Eastbank Fish Hatchery	Chinook	OtsEASTBK_sp	Spring	Interior stream type	*	*	*	*	*	X	X	X	X	X	X	
23	Eastbank Fish Hatchery	Chinook	OtsEASTBK_su	Summer	Interior ocean type	*	*	*	*	X	X	X	X	X	X	X	
24	Entiat National Fish Hatchery	Chinook	OtsENFH_su	Summer	Interior ocean type	*	*	*	*	*	X	X	X	X	X	X	
4	Kalama Falls	Chinook	OtsKAL_sp	Spring	Interior stream type	*	*	*	*	*	*	*	X	X	X	**	
4	Kalama Falls	Chinook	OtsKAL_fa	Fall	Interior stream type	*	*	*	*	*	*	*	*	X	X	**	
12	Klickitat State Fish Hatchery	Chinook	OtsKH_sp	Spring	Interior stream type	X	X	X	X	X	X	X	X	X	X	X	
12	Klickitat State Fish Hatchery	Chinook	OtsKH_fa c	Fall	Interior ocean type	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	X	
22	Leavenworth National Fish Hatchery	Chinook	OtsLNFH_sp	Spring	Interior stream type	*	*	*	*	*	X	X	X	X	X	X	
5	Speelyai Hatchery	Chinook	OtsLEW_sp	Spring	Interior stream type	*	*	*	*	*	*	*	X	X	X	**	
9	Little White Salmon National Fish Hatchery	Chinook	OtsLWS_fa	Fall	Interior ocean type	*	*	*	*	*	X	X	X	X	X	X	
9	Little White Salmon National Fish Hatchery	Chinook	OtsLWS_sp	Spring	Interior stream type	*	*	*	*	*	X	X	X	X	X	X	
9	Little White Salmon National Fish Hatchery - Touchet River	Chinook	OtsTOUCH_sp	Spring	Interior stream type	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	X	
42	Marion Forks Hatchery	Chinook	OtsNSANT_sp	Spring	Interior stream type	*	*	*	*	*	*	*	*	*	*	X	
27	Methow State Fish Hatchery	Chinook	OtsMETH_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
10	Parkdale Fish Facility	Chinook	OtsPFF_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
20	Priest Rapids Hatchery	Chinook	OtsPRH_fa	Fall	Interior ocean type	*	*	*	*	X	X	X	X	X	X	X	
14	Round Butte Fish Hatchery	Chinook	OtsRB_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
19	Ringold Springs State Hatchery	Chinook	OtsRG_fa	Fall	Interior ocean type	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	X	X	X	
41	South Santiam Hatchery	Chinook	OtsSSANT_sp	Spring	Interior stream type	*	*	*	*	*	*	*	X	X	*	X	
11	Spring Creek National Fish Hatchery	Chinook	OtsSPCR_fa	Fall	Lower Columbia	*	*	*	*	*	*	*	X	X	X	X	
3	North Toutle Hatchery	Chinook	OtsTOU_fa	Fall	Interior ocean type	*	*	*	*	*	*	*	X	X	X	**	
15	Three mile dam, Umatilla River	Chinook	OtsUMA_fa b	Fall	Interior ocean type	*	*	*	*	X	X	X	~	~	~	X	
17	South Fork Walla Walla facility	Chinook	OtsUMA_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
6	Washougal	Chinook	OtsWAS_fa	Fall	Interior ocean type	*	*	*	*	*	*	*	X	X	X	**	
13	Warm Springs National Fish Hatchery	Chinook	OtsWSNFH_sp h	Spring	Interior stream type	*	*	*	*	X	X	X	~	~	~	X	
25	Wells Fish Hatchery	Chinook	OtsWELLS_su	Summer	Interior ocean type	*	*	*	*	X	X	X	X	X	X	X	
28	Winthrop National Fish Hatchery	Chinook	OtsWTP_sp	Spring	Interior stream type	*	*	*	*	*	X	X	X	X	X	X	
18	Yakima Nation Prosser Hatchery	Chinook	OtsPRO_fa	Fall	Interior ocean type	*	*	*	*	X	X	*	X	X	X	X	
21	Levi George/Cle Elum (Integrated)	Chinook	OtsYRint_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	
21	Levi George/Cle Elum (Segregated)	Chinook	OtsYRseg_sp	Spring	Interior stream type	*	*	*	*	X	X	X	X	X	X	X	

Note: Species-specific collections code along with run type and genetic lineage are provided for both species. Map numbers correspond with Figure 13.

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- 337
- 338
- X
- X
- X
-
- a
- b
- c
- h
- N/A
- *
- **
- Chinook tissues genotyped using 343 SNPs
- Chinook tissues genotyped using 298 SNPs
- Chinook tissues genotyped using 96 SNPs
- Chinook broodstock sampled, spawned at another hatchery and genotyped using 298 SNPs
- Chinook Lyons Ferry stock consolidated under 'OtsLYON' starting in 2012
- Chinook Umatilla fall stock spawned at Little White Salmon Hatchery in 2015 & 2016; not distinguished from LWS stock. Broodstock collected/spawned at Ringold Springs in 2017; not distinguished from Ringold stock.
- Chinook typically spawned at Little White Salmon NFH, but due to low returns in 2018 they were spawned at Klickitat Hatchery.
- Chinook Warm Springs NFH spring stock spawned at Little White Salmon Hatchery starting in 2015-2017.
- Stock discontinued/non-existent
- Broodstock not sampled
- Broodstock sampled, tissues archived until funding identified for processing

Appendix 7. Checklist of PBT broodstock collections that comprise the PBT baselines for steelhead from 2008 through 2018.

Map num.	Hatchery	Species	Code	Run type	Lineage	Year											
						2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	
31	Dworshak National Fish Hatchery	Steelhead	OmyDWOR	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
35	Little Sheep Creek Hatchery	Steelhead	OmyLSCR	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery- Touchet	Steelhead	OmyTOUW c	Summer	Interior	*	X	X	X	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery	Steelhead	OmyLYON d	Summer	Interior	*	X	X	X	X	N/A	N/A	N/A	N/A	N/A	N/A	
29	Lyons Ferry Fish Hatchery - Grande Ronde	Steelhead	OmyCGRW c	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery - Tucannon	Steelhead	OmyTUCW c	Summer	Interior	*	X	X	X	X	X	X	X	X	X	X	
29	Lyons Ferry Fish Hatchery - Wallowa	Steelhead	OmyWALW	Summer	Interior	N/A	N/A	N/A	N/A	N/A	N/A	N/A	X	N/A	N/A	N/A	
37	Oxbow	Steelhead	OmyOXBO	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
39	Sawtooth Fish Hatchery	Steelhead	OmySAWT	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
39	Sawtooth Fish Hatchery - East Fork Salmon	Steelhead	OmyEFSW e	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
39	Sawtooth Fish Hatchery - Squaw Creek	Steelhead	OmySQUW f	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
40	Pahsimeroi Fish Hatchery	Steelhead	OmyPAHH	Summer	Interior	X	X	X	X	X	X	X	X	X	X	X	
34	Wallowa Fish Hatchery	Steelhead	OmyWALL	Summer	Interior	*	X	X	X	X	X	X	X	X	X	X	
23	Eastbank Hatchery	Steelhead	OmyEASTBK	Summer	Interior	*	*	*	*	X	X	X	X	X	X	X	
27	Methow Hatchery (Twisp)	Steelhead	OmyTWP i	Summer	Interior	*	*	*	*	*	X	X	X	X	~	~	
10	Parkdale Fish Facility	Steelhead	OmyPFF	Winter	Coastal	*	*	*	*	X	X	X	X	X	X	X	
14	Round Butte Fish Hatchery	Steelhead	OmyRB	Summer	Interior	*	*	*	*	*	X	X	X	X	X	X	
7	Skamania Hatchery (Summer)	Steelhead	OmySKH_su g	Summer	Coastal	*	*	*	*	*	X	X	X	X	X	X	
7	Skamania Hatchery (Winter)	Steelhead	OmySKH_wi g	Winter	Coastal	*	*	*	*	*	X	X	X	X	X	X	
16	Minthorn Springs	Steelhead	OmyUMA	Summer	Interior	*	*	*	*	X	X	X	X	X	X	X	
25	Wells Hatchery - On Station	Steelhead	Omy_WEL	Summer	Interior	*	*	*	*	*	X	X	X	X	X	X	
25	Wells Hatchery - Okanogan stock	Steelhead	OmyWEL_OKA	Summer	Interior	*	*	*	*	*	X	X	X	X	X	X	
25	Wells Hatchery - Omak stock	Steelhead	OmyWEL_OMA	Summer	Interior	*	*	*	*	*	X	X	X	X	X	X	
25	Wells Hatchery - Methow stock	Steelhead	OmyWEL_MET	Summer	Interior	*	*	*	*	*	X	X	X	X	X	X	
28	Winthrop National Fish Hatchery	Steelhead	OmyWTP	Summer	Interior	*	*	*	*	X	X	X	X	X	X	X	

- 340
- Note map numbers correspond to sites in Figure 13.
- 341
- X Steelhead tissues genotyped using 96 SNPs
- 342
- X Steelhead tissues genotyped using 379 SNPs
- 343
- X Steelhead tissues genotyped using 269 SNPs
- 344
- X Steelhead tissues genotyped using 192 SNPs
- 345
- X Steelhead tissues genotyped using 390 SNPs
- 346
- X Steelhead tissues genotyped using 368 SNPs
- 347
- ~ Steelhead broodstock sampled, spawned at another hatchery and genotyped using 379 SNPs
- 348
- Steelhead broodstock sampled, spawned at another hatchery and genotyped using 368 SNPs
- 349
- c Steelhead Lyons Ferry stock consolidated under 'OmyLYON' starting in 2012
- 350
- d Steelhead Lyons Ferry stock discontinued starting in 2013
- 351
- e Steelhead Sawtooth stock consolidated under 'OmySAWT' from 2012-2013
- 352
- f Steelhead Sawtooth stock consolidated under 'OmySAWT' in 2012; renamed 'Upper Salmon B-run' (YFLW) and consolidated under 'OmyPAHH' starting in 2013; spawned at Yankee Fork Weir by Shoshone-Bannock tribe beginning in 2017
- 353
- g Steelhead Skamania stock is collected late in calendar year, and is designated for the following broodyear (i.e., late 2012 collections are part of BY2013). Skamania winter Steelhead stock changed from Chambers Creek stock to Big Creek stock starting with SY2018.
- 354
- i Steelhead Methow Hatchery Twisp stock spawned at Winthrop NFH starting in 2017; not distiguished from Withrop stock
- 357
- N/A Stock discontinued/non-existent
- 358
- * Broodstock not sampled
- 359
- ** Broodstock sampled, tissues archived until funding identified for processing

Map num.	Hatchery	Species	Code	Run type	Lineage	Year										
						2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018
15	Three mile dam, Umatilla River	Coho	OkiUMA	Fall	Lower Columbia								X	X	X	X
22	Leavenworth National Fish Hatchery	Coho	OkiLNFH	Fall	Lower Columbia				X	X	X	X	X	X	X	X
30	Nez Perce Tribal Fish Hatchery	Coho	OkiNPT	Fall	Lower Columbia						*	*	X	X	X	X
18	Yakima Nation Prosser Hatchery	Coho	OkiPRO	Fall	Lower Columbia									X	X	X
28	Winthrop National Fish Hatchery	Coho	OkiWTP, OkiMET	Fall	Lower Columbia					X	X	X	X	X	X	X
33	Lookingglass Fish Hatchery - Lostine River	Coho	OkiLSTW	Fall	Lower Columbia											X

Note: map numbers correspond to sites in Figure 13.
X Coho tissues genotyped at 257 loci
* Samples received, but not genotyped.

Section 3: Genetic Stock Identification of Chinook Salmon, Sockeye Salmon, and Steelhead Harvest Mixtures in the Mainstem Columbia River

Introduction

Genetic Stock Identification (GSI) methods have proven to be effective in determining the proportion of stock origin in mixed stock applications of salmonids (Shaklee et al. 1999, Winans et al. 2004, Beacham et al. 2006, and Beacham et al. 2011). These methods have been demonstrated to be useful even at relatively fine geographic scales within the Columbia River Basin (CRB) (Hess et al. 2011, Hess and Narum 2011, Hess et al. 2014). Within the CRB, Chinook salmon consist of three major genetic lineages and steelhead consist of two major genetic lineages that can be further divided into populations that are genetically structured on a finer spatial scale (e.g., Waples et al. 2004; Narum et al. 2010; Blankenship et al. 2011). In this study, we used separate groups of SNP markers to discriminate 19 reporting groups for Chinook salmon, 14 reporting groups for steelhead, and four reporting groups for sockeye salmon.

Despite continuous improvements of the power of our Chinook salmon and steelhead baselines in GSI applications (Hess et al. 2014), we have determined that further improvement in the detail of data and accuracy of stock assignments could be made by utilizing a recently developed genetic technology (i.e., parentage based tagging (PBT)), in combination with GSI, in a tiered approach for stock identification (Hess et al. 2016). PBT is an efficient approach for mass tagging of fish. The method is carried out by first genotyping a set of potential parents which then provides the opportunity to assign a set of genotyped offspring to their true parent pair. PBT is currently being utilized on a broad scale in the Columbia River Basin, and was recently (i.e., 2012-present) expanded beyond Snake River hatcheries (Steele et al. 2011) to tag all Chinook salmon and steelhead hatchery broodstock from hatcheries in the CRB above Bonneville Dam. This application has effectively tagged all Snake River hatchery Chinook salmon and steelhead starting with the 2008 brood years, and elsewhere in the CRB above Bonneville Dam beginning with the 2012 brood year. When parent pairs of hatchery fish are identified with PBT, we can provide accurate information including age of the fish and the source hatchery in which its parents were spawned (Steele et al. 2011). We use PBT in this harvest study to identify hatchery-origin fish, and then use GSI to estimate stock-of-origin of all other hatchery fish that were not assigned with PBT and for all natural-origin fish. For sockeye salmon, we no longer rely solely on GSI to determine stock of origin because PBT can be used to identify reintroduced fish to the Yakima River. For the 2018 Chinook harvest, multiple age classes (3-, 4-, and 5-year old fish) can be identified from Lower Columbia, Middle Columbia, and Snake River stocks using PBT (Appendix 6).

We continue to employ the genotyping-in-thousands by sequencing (GT-seq) approach that has been developed in our laboratory (Campbell et al. 2015). This approach has increased the cost-effectiveness for genotyping moderate numbers of SNP loci (100s) for relatively large numbers of individuals (1000s), which allows us to run all SNP loci regardless of whether we intend to use primarily PBT analyses or a combination of PBT and GSI. Thus, our projects now benefit from the additional data that comes from genotyping with all available markers (i.e., increased power for statistical assignment of individuals).

Fisheries conducted in the mainstem of the lower and middle Columbia River provide an important application of genetic stock analyses because the fish harvested consist of mixtures of stocks throughout the CRB. Further, mainstem Chinook and Sockeye salmon fisheries represent

a majority of the CRB harvest of this species taken by the commercial, sport, and tribal fishermen. In order to help support sustainable fisheries, PBT and GSI can be used to address two primary questions: 1) how are Chinook salmon stocks temporally and spatially distributed in the mainstem Columbia River; and 2) how are these stocks temporally and spatially distributed in the harvests of fisheries. Importantly, we can now estimate abundance of all genetic stocks using the reported catch estimates of each fishery.

Project objectives and higher-level harvest management questions

Our study had three primary objectives: 1) utilize a combination of PBT and GSI analyses to determine stock composition of Chinook salmon harvested in sport, commercial, and tribal fisheries in the mainstem Columbia River, 2) utilize GSI to estimate stock composition of sockeye salmon harvested above and below Bonneville Dam in commercial, sport, and tribal fisheries, 3) estimate abundance of all stocks based on catch estimates for each fishery. Results from these objectives were used to address:

Harvest RM&E: F&W Program Management Question: What are your in-river monitoring results and what are your estimates of stock composition and stock-specific abundance, escapement, catch, and age distribution?

Increasingly, we are tailoring our analyses to address specific questions that fisheries managers have presented to us. For example, in 2012 managers proposed extending the geographic boundary of one of the mark selective spring-run Chinook salmon sport fisheries above Bonneville Dam that occurs at the mouth of the Wind River. This extension created a larger “bubble” boundary at the mouth of the Wind River and was intended to increase Columbia River mainstem fishing access while maintaining targeted focus on Wind River spring-run Chinook salmon. For 2012-2016, we examined the stock composition of the Wind River sport harvest and provided context by comparing stock proportions among the various samples from other fisheries and Bonneville Dam that were analyzed that same year. The Wind River sport fishery has not been sampled since 2017 and could not be included in analyses. However, there is a new research gear (pound net) being tested by the joint states in the fall fisheries below Bonneville Dam. We have included analysis of the 2018 fall pound net fishery to estimate stock abundance of fall Chinook salmon among the clipped fish that were retained (kept) and the unclipped fish that were released. We included in this report the 6th year of analysis of Sockeye salmon fisheries in the Columbia River mainstem. Differences in relative abundance of the three main stocks (Okanagan, Wenatchee, and Snake) present challenges to managing lower river harvest, because of the desire to harvest the highly abundant Okanagan stock around the much less abundant Snake River stock and moderately abundant Wenatchee River stock. Stock composition estimates are expected to help determine how harvest is impacting these various stocks. This year, we identified a stock of reintroduced Sockeye salmon to the Yakima River using PBT and can now estimate the contribution of this reintroduced stock to each harvest of Sockeye salmon and the run at large.

Time line for completion of objectives

Objectives will be ongoing and PBT/GSI results updated each year for harvest analyses of salmonids throughout the accords-funding. As new genetic techniques are developed, they will be applied to this project and results will be compared between years to determine the extent of improvements.

Our study was not designed to address the following question:

Harvest RM&E: F&W Program Management Question: Can selective fisheries targeting hatchery fish or healthy populations reduce impacts on ESA-listed populations?

Accuracy testing of PBT and GSI baselines

Prior to conducting analyses for fisheries harvest collections and mixture samples encountered at Bonneville Dam (Section 4), we assessed the accuracy of our PBT and GSI baselines in assigning Chinook salmon and steelhead to their hatchery brood or reporting group of origin (see Results section).

Methods

Methods for estimating stock composition are available at (<https://www.monitoringmethods.org/Protocol/Details/229>). The Monitoring Methods Protocol is entitled Snake River steelhead and Chinook salmon stock composition estimates (2010-026-00) v1.0.

Tissue collection of Chinook salmon and Sockeye salmon

Tissues were sampled from Chinook salmon in 2018 from a total of 9 different mixture sources: the spring-run seasons of the following fisheries: 1) lower river test, 2) lower river sport, and 3) Zone 6 Treaty ceremonial permit, the summer management period harvests of the following fisheries: 4) lower river sport, and 5) Zone 6 tribal summer, and the fall-run harvest from 6) the lower river test (pound net), 7) lower river commercial, 8) lower river sport, and 9) Zone 6 tribal fall fishery. Tissues are also collected from steelhead, Chinook salmon, and Sockeye salmon at Bonneville Dam for stock ID (see Section 4). While fisheries generally harvest jack sized Chinook salmon at low rates and do not have specific harvest limits on jacks, jacks do comprise part of the harvest and may be sampled if encountered. Jacks are sampled at the Bonneville AFF trap in the proportion that they are encountered in the sampling. Sampling restrictions at the AFF can result in biases in the size of fish sampled compared to the run at large. Therefore, we split the AFF sample of Chinook salmon into adults and jacks and analyze them separately. Harvest tissues were collected in coordination with existing monitoring programs led by Washington Department of Fish and Wildlife (WDFW) and Oregon Department of Fish and Wildlife (ODFW) and the Yakama Nation. The spring management period Chinook salmon fisheries were sampled below Bonneville Dam in the sport, test fishery, and in the Treaty permit fishery (Figure 12; Table 6). The summer management period fisheries were sampled below Bonneville Dam in the sport fishery, and above Bonneville Dam in the Zone 6 Treaty commercial fishery. Due to limited funds, we analyzed a subset of samples obtained from the various fisheries sampled above and below Bonneville Dam. For all fisheries in which we had to subsample the harvest, we selected fish randomly and with a balanced design across spatial regions. However, each subsample should be tailored to the way the catch is reported for each fishery to accurately represent the geographic and temporal stratification of the CREEL.

Stock proportions were estimated and expanded by the catch reported for each fishery source, such that stock abundance could be compared across fisheries. We use the following four main geographic regions (Figure 12): Region A corresponds to our grouping of pre-existing Oregon and Washington state sport fishing zones 1-4 (or commercial zones 4-5), Region B corresponds to our grouping of sport zones 5-10 (or commercial zones 1-3). Here, we do not discriminate between Region 01 and Region 02 in the Zone 6 fishery, because that information did not accompany the samples we received. These sets of groupings were established for this study in order to achieve balanced sampling for analysis of these fishery datasets, as well as to best match the stratification of the reported catch.

Non-Treaty fisheries during the spring management period for Chinook salmon are mark-selective based on absence or presence of the adipose fin to distinguish hatchery fish from natural origin fish, respectively. These adipose markings make it possible to have a mark-selective sport and commercial fishery in which only fish with missing adipose fins (hatchery-origin) are legally retained. Fish with intact adipose fins that are caught in these fisheries are released, but mortality rates are unknown from these releases. In addition to sampling hatchery-origin fish from the mark selective commercial and sport fisheries, we were able to obtain samples from non-clipped hatchery and natural origin fish from Bonneville Dam and the Treaty Zone 6 fishery above Bonneville Dam.

Tissues were sampled from sockeye salmon in 2018 from three fishery mixture sources: 1) lower river sport, 2) Bonneville Dam (see Section 4), and 3) the Treaty fishery in Zone 6. All samples obtained from these fisheries were analyzed.

Molecular data

Methods for DNA extraction, DNA amplification, and genotyping of SNP assays using genotyping-in-thousands by sequencing (GT-seq) are available at (<https://www.monitoringresources.org/Document/Method/Details/5446>). Additional details regarding how 192 SNPs were reduced to 186 SNPs can be found in Hess et al. (2012, 2013). Subsequently, we have reduced our Chinook salmon GSI baseline from 186 SNPs to 179 SNPs (Hess et al. 2019), and further reduced from 179 SNPs to 177 SNPs because we were unable to transition the full set of 186 SNPs to GT-seq protocols and subsequent iterations of our active GT-seq panels. These 177 SNP markers were used for GSI, and for PBT analyses, we used 93 of the SNPs (legacy panel) as well as an expanded panel of 254 SNPs. We used 363 SNP markers for GSI and PBT of sockeye mixtures.

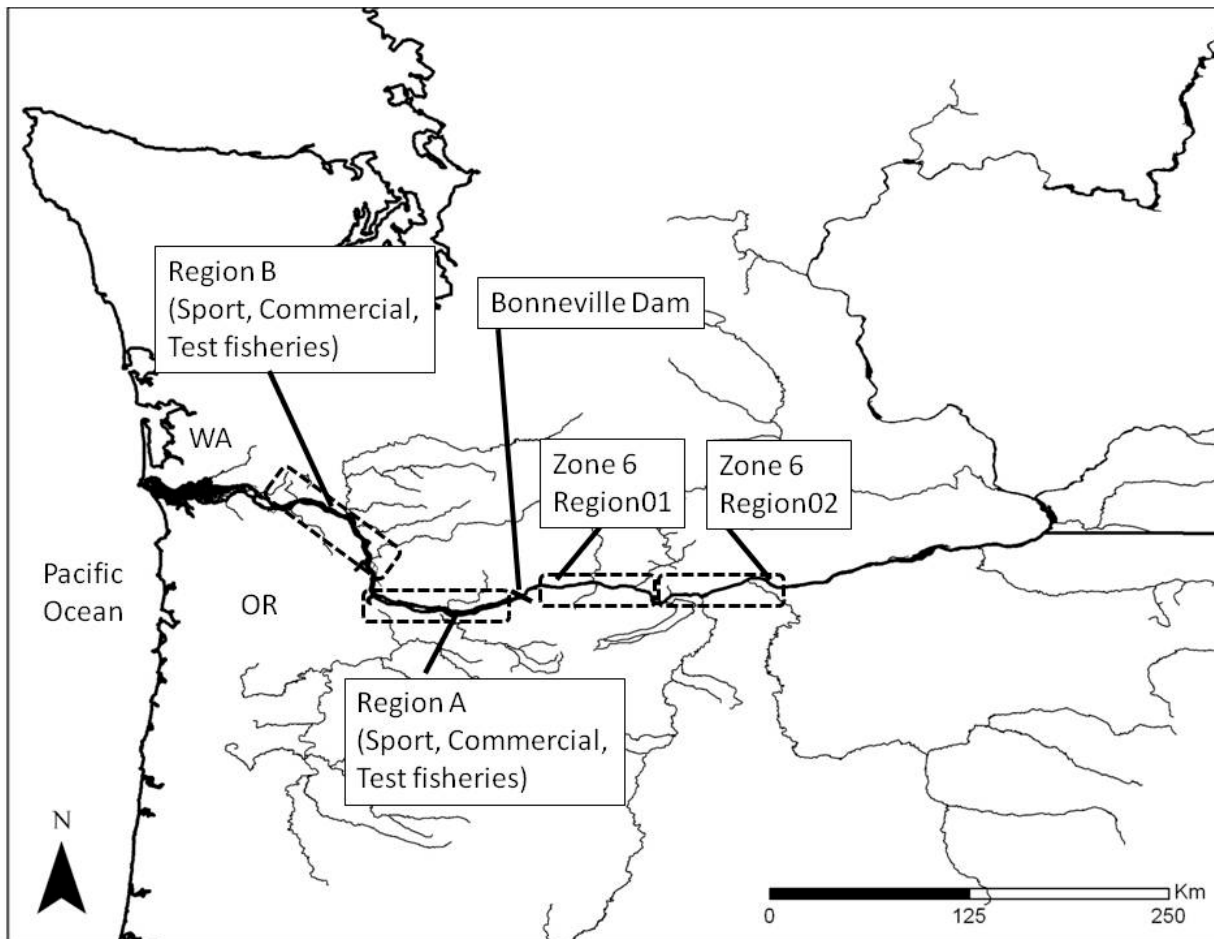


Figure 12. Project scope showing sources of Chinook salmon and Sockeye salmon harvest mixtures that were analyzed using PBT/GSI.

537 **Table 6. Characteristics of Chinook salmon harvest samples by fishery, region, and adipose-clip status by weekly strata in 2018.**

						Spring										Summer										Fall										
						Statistical weeks																														
Period	Fishery	Region	Clip	Sampled	Genotyped	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40
Spring	Sport	A	AD	354	347		8	15	86	125	44						11	27	9	22																
		B	AD	527	526	1	12	46	62	300	35							7	8	16	39															
	Test	B	AD	571	567		2	5	15	37	78	154	191	82	3																					
			AI	86	83			1	7	2	20	21	26	6																						
	Ceremonial	Below_BON	AD	240	235									26	165	44																				
			AI	39	38									2	30	6																				
		ZN6	AD	139	139									6	43	60	30																			
			AI	29	29									2	10	8	9																			
Summer	Sport	A	AD	31	30															7	22					1										
		B	AD	43	42															11	28	2				1										
	Treaty	ZN6	AD	166	165															66	41	30	23	3	2											
			AI	59	58															24	15	13	3		3											
Fall	Commercial	A	AD	291	198																									87	111					
			AI	240	100																									50	50					
	Sport	A	AD	45	45																							2	3		3	23	14			
			AI	174	119																							1	2	2	16	49	49			
		B	AD	33	32																							2	3	10	13	4				
			AI	80	79																							1	17	25	29	6	1			
	Test	B	AD	646	299																									1	190	53	53	2		
			AI	1050	172																									21	88	28	33	2		
	Treaty	ZN6	AD	370	234																							2		2	25	50	66	32	42	15
			AI	1387	456																								3	6	1	53	51	100	89	99
				6600	3993	1	22	67	170	464	177	175	253	336	121	39	18	35	25	61	108	106	45	26	3	6	1	11	31	199	578	264	316	125	141	69

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PBT Rate Expansion using SCOBIDEUX and SPIBETR functions

The use of PBT adds complexity to analyses when each hatchery broodstock is genotyped at rates less than 100%. Expansion of hatchery-origin adipose-intact fish can lead to bias when we characterize natural-origin stocks by attributes (Figure 13). We have developed and implemented a fully automated method which minimizes the bias that PBT rate expansion can impose (Delomas and Hess, In review). The correction implemented by this method (SPIBETR, Salmonid Prior Information to Balance Expansion from Tag Rates) is illustrated below:

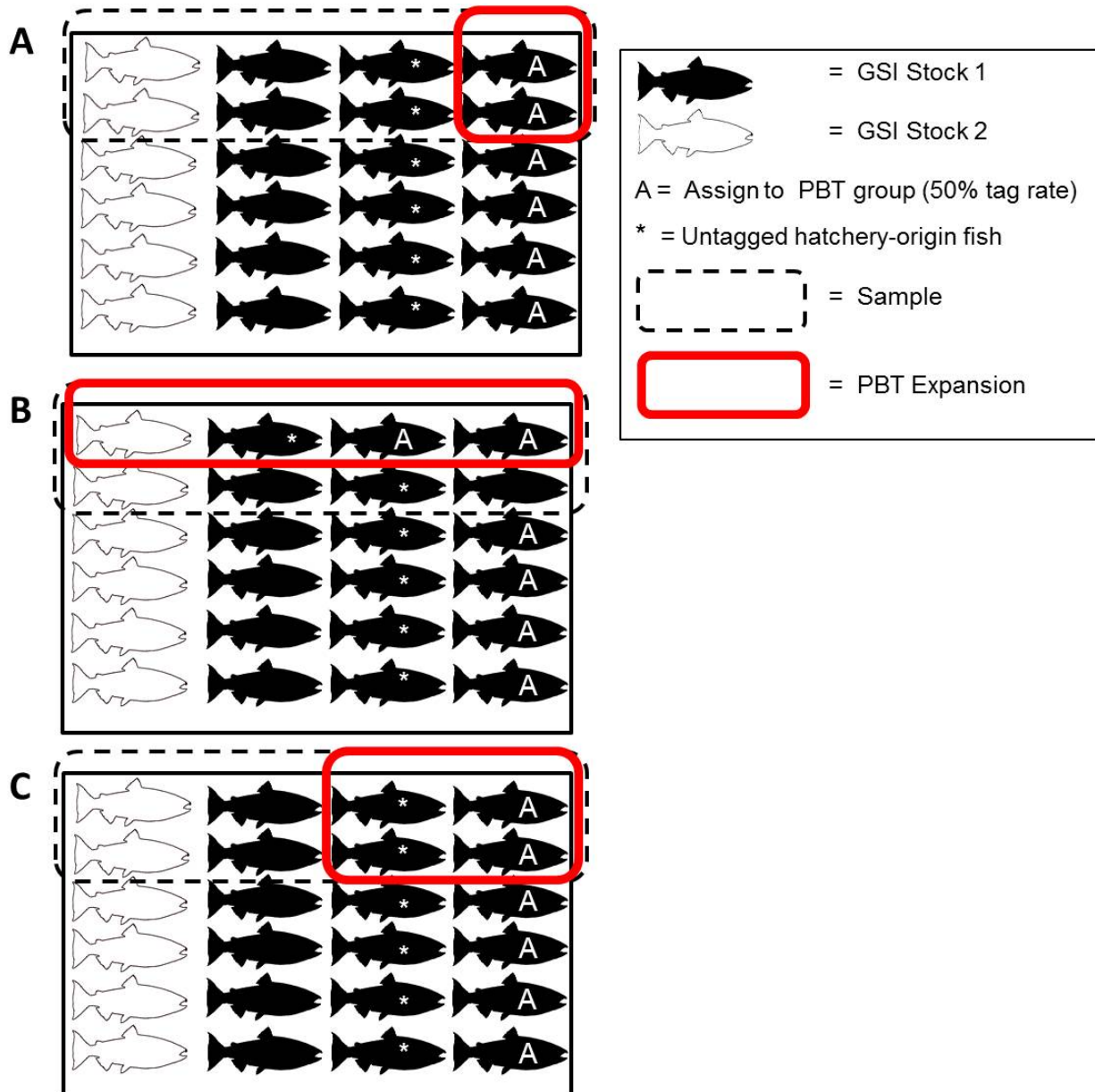


Figure 13. Conceptual illustration of the bias correction the SCOBIDEUX method provides.

In **Figure 13**, each pane shows a mixture of fish consisting of two natural-origin stocks (GSI Stock 1 and 2) and one PBT group with a 50% tag rate. The true proportions are 50% hatchery

origin, 25% natural-origin GSI stock 1, and 25% natural-origin GSI stock 2. When PBT rates are ignored, we would estimate the mixture to be 25% hatchery origin, 50% natural-origin GSI stock 1, and 25% natural-origin GSI stock 2 (**Figure 13, A**); this estimate is biased low toward the hatchery-origin fish. However, if we were to take PBT rates into account but did not adjust for the GSI information in the mixture, we could end up with estimates that accurately reflect the true proportion of hatchery-origin fish but do not accurately reflect the true stock composition of the natural-origin fish: e.g., **Figure 13, B**: yields an estimate of 50% hatchery origin, 37.5% natural-origin GSI stock 1, and 12.5% natural-origin GSI stock 2. Finally, using the corrections implemented by the SCOBIDEUX and SPIBETR functions, we are able to expand GSI stock 1 by PBT rates and correct this expansion by proportionally decreasing the GSI stock 1 in the remaining sample mixture, which yields scenario C (**Figure 13**): the mixture would be estimated to be 50% hatchery origin, 25% natural-origin GSI stock 1, and 25% natural-origin GSI stock 2.

GSI baselines for Chinook salmon, sockeye salmon, and steelhead

Chinook salmon GSI analyses were performed using the updated baseline referred to as “Columbia River Basin Chinook salmon GSI baseline version 3.1” and is available on the FishGen website (<https://www.fishgen.net>). However, we made on slight modification and decreased the number of SNP markers to 177 that were included in the latest iteration of the GT-seq panel. This baseline consists of 61 collections that are delineated into the following 19 reporting groups: Columbia Rogue “01_YOUNGS”, West Cascade spring-run “02_WCASSP”, West Cascade fall-run “03_WCASFA”, Willamette River spring-run “04_WILLAM”, Spring Creek Group Tule fall-run “05_SPCRTU”, Klickitat River spring-run “06_KLICKR”, Deschutes River spring-run “07_DESCSP”, John Day River spring-run “08_JOHNDR”, Yakima River spring-run “09_YAKIMA”, upper Columbia River spring-run “10_UCOLSP”, Tucannon River spring-run “11_TUCANO”, Hells Canyon spring-run “12_HELLSC”, South Fork Salmon River spring-run “13_SFSALM”, Chamberlain Creek spring-run “14_CHMBLN”, Middle Fork Salmon River spring-run “15_MFSALM”, upper Salmon River spring-run “16_UPSALM”, Deschutes River fall-run “17_DESCFA”, upper Columbia River summer-/fall-run “18_UCOLSF”, and Snake River fall-run “19_SRFALL” (Table 7; Figure 14). Reporting groups were primarily determined by the relative genetic similarity among populations according to a phylogenetic analysis, and our previous results demonstrate sufficient power to discern three reporting groups (17_DESCFA, 18_UCOLSF, and 19_SRFALL) among the interior ocean-type collections. In one year, we had grouped all interior ocean-type collections into a single reporting group “Interior_Columbia_R_su/fa” (Hess et al. 2013). Genetic distances were computed from allele frequencies based on Nei’s (1972) genetic distance, with the PHYLIP v 3.69 (Felsenstein 1989) and 1000 bootstrap replicates were performed. Distances were clustered using the Neighbor – Joining method (Saitou and Nei, 1987), and a consensus tree was constructed (<http://evolution.genetics.washington.edu/phylip/>) (Figure 15). The clusters are labeled with names of reporting groups used to aggregate the collections based on a combination of factors including genetic similarity, life history, and geographic proximity. Bootstrap support is shown with shaded ovals (Source: Hess et al. 2015).

The 10_UCOLSP reporting group includes the following Bonneville pool hatchery stocks: Carson stock (Ots22), and Little White Salmon R. (Ots23) because they are genetically indistinguishable from Upper Columbia R. spring Chinook salmon (includes Walla Walla and

Umatilla River stocks). This composite group is notable because inclusion of these Bonneville pool stocks explains why a large proportion of fish from the Wind R. sport fishery should assign to this 10_UCOLSP reporting group. However, the PBT baseline is now able to specifically identify fish from Carson Hatchery and Little White Salmon Hatchery allowing them to be distinguished from fish returning to the upper Columbia R. The 01_YOUNGS reporting group represents an out-of-basin genetic stock (originating from the Rogue R., OR) that is reared within the Columbia R. at Youngs Bay. Basic QAQC was performed to remove duplicate individuals and strays from the reference populations in the baseline. The baseline and reporting group data is available on FishGen.

GSI analyses for *O. nerka* utilized a baseline that included sockeye salmon and kokanee populations from throughout the Columbia River basin. This baseline included sockeye populations from the Osoyoos (i.e., Okanogan), Wenatchee, and Redfish Lake (i.e., Snake), and a kokanee population from Lake Whatcom that were included in “Sockeye GSI baseline v1.0, and were shown to accurately discriminate among these major stock (Hess et al 2013). We updated our baseline to included additional kokanee populations from Alturas Lake, Fishhook Creek, Lake Billy Chinook, Meadow Creek, Suttle Creek, Cougar, Gold, North Fork Tieton, Odell, Speylai, Stanley, Warm, Wizard, Wallowa River, and Wallowa Lake, and refer to this as “Sockeye GSI baseline v3.0”. The transition to GT-seq required omission of a few loci due to poor genotyping quality with the new protocols. A total of 363 SNPs was used for these analyses.

For steelhead, we have the following two GSI baselines available: 1) GSI baseline version 3.3 with 177 SNPs and 2) a new GSI baseline with 335 SNPs. The GSI version 3.3 comprises 116 collections from throughout the Columbia River basin that are partitioned into the following 14 reporting groups: 01_WCOAST (Quinault River), 02_LOWCOL (lower Columbia River), 03_SKAMAN (Skamania hatchery releases at three sites in lower Columbia River, Willamette River, and Klickitat River), 04_WILLAM (Willamette River), 05_BWSALM (Big White Salmon River), 06_KLICKR (Klickitat River), 07_MGILCS (middle Columbia River, Grande Ronde River, Imnaha River, lower Snake River, lower Clearwater River, and lower Salmon River), 08_YAKIMA (Yakima River), 09_UPPCOL (upper Columbia River), 10_SFCLWR (South Fork Clearwater River), 11_UPCLWR (upper Clearwater River), 12_SFSALM (South Fork Salmon River), 13_MFSALM (Middle Fork Salmon River), and 14_UPSALM (upper Salmon River) (see Table 8; Figure 16). Genetic distances were computed from allele frequencies based on Nei’s (1972) genetic distance, with the PHYLIP v 3.69 (Felsenstein 1989) and 1000 bootstrap replicates were performed. Distances were clustered using the Neighbor – Joining method (Saitou and Nei, 1987), and a consensus tree was constructed (<http://evolution.genetics.washington.edu/phylip/>) (Hess et al. 2019). The GSI baseline with 335 SNPs comprises 128 collections throughout the Columbia River that are partitioned into 13 reporting groups (**Table 9**), which are all the reporting groups listed in version 3.3 except 01_WCOAST (Quinault River). This 335 SNP baseline is still in the testing phase and will be considered for exclusive use of GSI applications only if it is determined to be more accurate than the existing version 3.3. baseline.

641 **Table 7. Sample sizes and reporting groups of Chinook salmon baseline populations. Lineages: ST (stream type), OT (ocean type), LC (Lower Columbia).**

ID	Collection	(n)	Lineage	Reporting Groups	Reporting Group description
OTS01	Youngs Bay fall-run	91	Rogue	01_YOUNGS	Youngs Bay- Columbia Rogue stock
OTS02	Cowlitz R spring-run	90	LC	02_WCASSP	West Cascade spring-run
OTS03	Kalama R spring-run	83	LC	02_WCASSP	West Cascade spring-run
OTS04	Cowlitz R fall-run	82	LC	03_WCASFA	West Cascade fall-run
OTS05	Elochoman R fall-run	86	LC	03_WCASFA	West Cascade fall-run
OTS06	Lewis R fall-run	93	LC	03_WCASFA	West Cascade fall-run
OTS07	NF Lewis fall-run	178	LC	03_WCASFA	West Cascade fall-run
OTS08	Sandy R fall-run	83	LC	03_WCASFA	West Cascade fall-run
OTS09	McKenzie R spring-run	78	LC	04_WILLAM	Willamette River spring-run
OTS10	N Santiam R spring-run	79	LC	04_WILLAM	Willamette River spring-run
OTS11	Sandy R spring-run	48	LC	04_WILLAM	Willamette River spring-run
OTS12	White Salmon fall-run	77	LC	05_SPCRTU	Spring Creek tule fall-run
OTS13	Spring Creek NFH tule fall-run	49	LC	05_SPCRTU	Spring Creek tule fall-run
OTS14	Klickitat R spring-run	84	ST	06_KLICKR	Klickitat River spring-run
OTS15	Shitike R spring-run	93	ST	07_DESCSP	Deschutes River spring-run
OTS16	Warm Springs R spring-run	90	ST	07_DESCSP	Deschutes River spring-run
OTS17	John Day R spring-run	78	ST	08_JOHNDR	John Day River spring-run
OTS18	Middle Fork John Day R spring-run	47	ST	08_JOHNDR	John Day River spring-run
OTS19	North Fork John Day R spring-run	42	ST	08_JOHNDR	John Day River spring-run
OTS20	American R spring-run	76	ST	09_YAKIMA	Yakima River spring-run
OTS21	Cle-Elum spring-run	88	ST	09_YAKIMA	Yakima River spring-run
OTS22	Winthrop NFH spring-run	82	ST	10_UCOLSP	upper Columbia River spring-run/ Carson Hatchery spring-run
OTS23	little White Salmon R spring-run	93	ST	10_UCOLSP	upper Columbia River spring-run/ Carson Hatchery spring-run
OTS24	Wenatchee R spring-run	109	ST	10_UCOLSP	upper Columbia River spring-run/ Carson Hatchery spring-run
OTS25	Entiat R spring-run	98	ST	10_UCOLSP	upper Columbia River spring-run/ Carson Hatchery spring-run
OTS26	Tucannon R spring-run	81	ST	11_TUCANO	Tucannon River spring-run
OTS27	Wenaha R spring-run	179	ST	12_HELLSC	Hells Canyon spring-run
OTS28	Lostine R spring-run	212	ST	12_HELLSC	Hells Canyon spring-run
OTS29	Grande Ronde R spring-run	314	ST	12_HELLSC	Hells Canyon spring-run
OTS30	Imnaha R spring-run	96	ST	12_HELLSC	Hells Canyon spring-run
OTS31	Lolo Cr spring-run	89	ST	12_HELLSC	Hells Canyon spring-run
OTS32	Red R spring-run	221	ST	12_HELLSC	Hells Canyon spring-run
OTS33	Powell R spring-run	56	ST	12_HELLSC	Hells Canyon spring-run
OTS34	Red R weir spring-run	91	ST	12_HELLSC	Hells Canyon spring-run
OTS35	South Forth Salmon R spring-run	139	ST	13_SFSALM	South Fork Salmon River spring/summer-run
OTS36	Johnson Cr spring-run	137	ST	13_SFSALM	South Fork Salmon River spring/summer-run
OTS37	Secesh R spring-run	252	ST	13_SFSALM	South Fork Salmon River spring/summer-run
OTS38	Chamberlain Cr spring-run	219	ST	14_CHMBLN	Chamberlain Creek spring/summer-run
OTS39	Big Cr spring-run	139	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS40	Camas Cr spring-run	55	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS41	Loon Cr spring-run	107	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS42	Sulphur Cr spring-run	94	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS43	Bear Valley Cr spring-run	135	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS44	Capehorn Cr spring-run	214	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS45	Marsh Cr spring-run	228	ST	15_MFSALM	Middle Fork Salmon River spring/summer-run
OTS46	North Fork Salmon R spring-run	55	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS47	Lemhi R spring-run	96	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS48	Pahsimeroi R spring-run	92	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS49	East Fork Salmon R spring-run	286	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS50	Salmon R spring-run	83	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS51	West Fork Yankee Fork spring-run	75	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS52	Valley Cr spring-run	100	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS53	Sawtooth Hatchery weir spring-run	186	ST	16_UPSALM	upper Salmon River spring/summer-run
OTS54	upper Deschutes R fall-run	252	OT	17_DESCFA	Deschutes River fall-run
OTS55	lower Yakima R fall-run	62	OT	18_UCOLSF	upper Columbia River summer/fall-run
OTS56	Hanford Reach fall-run	93	OT	18_UCOLSF	upper Columbia River summer/fall-run
OTS57	Wenatchee R summer-run	92	OT	18_UCOLSF	upper Columbia River summer/fall-run
OTS58	Entiat R summer-run	51	OT	18_UCOLSF	upper Columbia River summer/fall-run
OTS59	Methow R summer-run	87	OT	18_UCOLSF	upper Columbia River summer/fall-run
OTS60	Lyons Ferry weir fall-run	90	OT	19_SRFALL	Snake River fall-run
OTS61	Clearwater R fall-run	228	OT	19_SRFALL	Snake River fall-run

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671 **Table 8. Results of the leave-one-out analysis for the steelhead GSI 335 SNP baseline.**

Reporting group	02_LOWCOL	03_SKAMAN	04_WILLAM	05_BWSALM	06_KLICKR	07.05_FIFTNM	07.1_DESCH	07.2_JDAY	07.3_UMAT	07.4_LOWSNAKE	07.5_IMNA	07.52_GrandeRonde	07.6_LOWCLWR	07.7_LOWSALM	08_YAKIMA	09_UPPCOL	10_SFCLWR	11_UPCLWR	12_SFSALM	13_MFSALM	14_UPSALM	Self	MCGILCS
02_LOWCOL	95.9	3.4	0.4	0.1	0.2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	95.9	
03_SKAMAN	4.8	95.1	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	95.1	
04_WILLAM	2.7	4.6	92.7	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	92.7	
05_BWSALM	1.5	0.0	0.6	92.7	3.3	1.0	0.0	0.0	0.0	0.0	0.0	0.0	0.5	0.0	0.0	0.0	0.1	0.0	0.0	0.0	0.1	92.7	
06_KLICKR	1.3	4.3	0.0	1.2	83.5	4.5	0.8	0.0	0.2	1.4	0.0	0.3	0.1	0.1	0.6	0.8	0.2	0.0	0.0	0.0	0.6	83.5	
07.05_FIFTNM	0.7	0.0	0.1	1.1	5.7	49.2	7.4	0.7	2.9	9.9	4.0	5.1	1.6	1.7	1.4	1.8	0.0	0.8	0.0	1.6	4.1	49.2	82.6
07.1_DESCH	0.0	0.0	0.0	0.3	0.4	3.2	65.2	4.1	3.5	5.8	1.2	7.8	2.1	1.0	0.8	2.5	0.2	0.0	0.0	0.2	1.8	65.2	93.9
07.2_JDAY	0.0	0.0	0.0	0.0	0.0	0.8	4.1	58.9	9.0	7.4	3.1	11.0	2.0	0.4	0.8	1.1	0.2	0.0	0.2	0.0	0.8	58.9	96.8
07.3_UMAT	0.0	0.0	0.0	0.0	0.0	1.1	7.2	13.5	36.1	16.7	0.3	12.1	4.1	0.8	4.6	2.5	0.1	0.0	0.0	0.0	0.7	36.1	92.0
07.4_LOWSNAKE	0.0	0.0	0.0	0.0	0.0	1.8	3.8	4.0	8.0	46.5	1.9	11.9	4.9	2.7	1.6	8.1	0.4	0.1	0.0	0.1	4.2	46.5	85.5
07.5_IMNA	0.0	0.0	0.0	0.0	0.0	0.5	3.0	3.2	2.6	9.6	61.6	8.1	2.3	4.5	0.4	1.0	0.5	0.0	1.0	1.6	61.6	95.5	
07.52_GrandeRonde	0.0	0.0	0.0	0.0	0.0	1.2	5.9	8.5	6.0	10.9	2.0	56.8	2.4	1.3	0.3	3.1	0.0	0.1	0.0	0.3	1.2	56.8	95.0
07.6_LOWCLWR	0.0	0.0	0.0	0.0	0.0	0.6	2.8	1.5	3.7	9.3	1.1	9.2	67.0	0.5	0.3	2.1	0.7	0.6	0.0	0.1	0.5	67.0	95.7
07.7_LOWSALM	0.0	0.0	0.0	0.0	0.0	0.2	2.5	0.3	1.8	13.2	3.5	5.3	1.4	61.8	0.0	1.4	0.1	0.0	0.7	2.7	5.1	61.8	90.0
08_YAKIMA	0.0	0.0	0.0	0.0	0.2	0.6	0.8	1.2	1.1	1.3	0.1	1.1	0.4	0.1	92.0	0.9	0.0	0.0	0.0	0.0	0.1	92.0	
09_UPPCOL	0.0	0.0	0.0	0.4	0.6	2.0	2.4	1.4	2.1	15.9	0.9	6.7	2.3	1.4	2.3	59.4	0.0	0.0	0.0	0.1	2.2	59.4	
10_SFCLWR	0.0	0.0	0.0	0.0	0.0	0.0	0.0	1.0	0.0	0.0	0.0	0.0	1.7	0.0	0.0	0.0	88.0	8.9	0.0	0.0	0.3	88.0	
11_UPCLWR	0.0	0.0	0.0	0.0	0.0	0.2	0.1	0.0	0.0	0.1	0.0	0.0	0.6	0.0	0.0	0.0	3.7	94.9	0.0	0.0	0.4	94.9	
12_SFSALM	0.0	0.0	0.0	0.0	0.0	0.0	0.2	0.0	0.0	0.1	0.1	0.2	0.0	0.9	0.0	0.0	0.0	0.0	98.2	0.2	0.1	98.2	
13_MFSALM	0.0	0.0	0.0	0.0	0.0	0.2	0.3	0.3	0.0	2.1	0.5	0.4	0.0	1.3	0.0	0.1	0.0	0.0	0.0	91.9	2.8	91.9	
14_UPSALM	0.0	0.0	0.0	0.0	0.2	0.6	1.1	0.3	0.3	7.6	1.5	2.8	1.6	3.7	1.9	3.0	0.4	0.1	0.0	3.5	71.5	71.5	

672 **Note:** Reporting group, population ID, population name and samples size used in analysis are provided for each collection in the baseline. The number of correct assignments to reporting group for each population in the baseline are reported (gray shading) and are tallied to provide the number of correct assignments for the reporting group
673 overall (yellow shading). The average proportion of correct assignments to reporting group (across collections) is provided along the diagonal. MCGILCS has been challenging to discriminate smaller groups that comprise the large region. This baseline of 335 SNPs had limited ability (correct assignment <70%) to accurately resolve
674 smaller groups (groups with “07” notation) within MCGILCS. Therefore we also report the correct assignments to the MCGILCS group as a whole (far right column; “MCGILCS”).

675 **Table 9. Sample sizes, locations, and reporting groups of steelhead baseline populations for**
676 **the new panel of 335 SNPs.**

Reporting Group	Population	Name	Latitude	Longitude	N
02_LOWCOL	POP001.1	ElochomanR	46.226734	-123.330417	55
	POP001.2	AbernathyCr	46.225600	-123.148100	96
	POP002	CowlitzR	46.502611	-122.588094	131
	POP003	GermanyCr	46.190147	-123.123783	43
	POP004	GraysR	46.367836	-123.581335	40
	POP005	KalamaR_wi	46.032897	-122.870311	41
	POP006	MillCr	46.188517	-123.177297	40
	POP007	NFLewisR	45.956781	-122.555108	92
	POP008	StillCr	45.330870	-121.915750	13
03_SKAMAN	POP009	WashougalR	45.585126	-122.393031	49
	POP010	ClackamasR_su	45.241700	-122.281700	47
	POP011	KalamaR_su	46.032897	-122.870311	42
04_WILLAM	POP012	KlickitatRSkamania_su	45.818764	-121.235121	271
	POP013	ClackamasR_wi	45.317613	-122.413773	261
	POP014	EagleCr	45.351428	-122.384017	23
	POP015	NSantiamR	44.750042	-122.403872	32
	POP016	NFEagleCr	45.351428	-122.384017	25
05_BWSALM	POP017	SFSantiamR	44.415003	-122.673942	74
	POP019	BuckCr_above	45.799317	-121.484314	44
	POP020	BuckCr_below	45.799317	-121.484314	38
	POP022	LRattlesnakeCr	45.799317	-121.484314	39
	POP023	BigWhiteSalmonR_mainstem	45.799317	-121.484314	36
	POP024	MidRattlesnakeCr	45.799317	-121.484314	31
	POP025	BigWhiteSalmonR_above	45.799317	-121.484314	24
	POP026	RattlesnakeCr	45.799317	-121.484314	41
	POP028	BigWhiteSalmonR_Husum	45.799550	-121.485590	14
06_KLICKR	POP031	BowmanCr	45.845210	-121.042110	99
	POP032	DeadCanyonCr	45.942280	-121.143940	34
	POP033	DillacortCr	45.741407	-121.222239	26
	POP034	WheelerCr	45.795093	-121.192829	8
	POP035	SnyderCr	45.824664	-121.156523	29
	POP036	SwaleCr	45.809090	-121.065160	91
	POP037	LKlickitatR	45.841340	-121.003888	77
	POP038	LowSummitCr	45.935701	-121.101827	64
	POP039	LowTroutCr	46.037760	-121.199420	74
	POP040	LowWhiteCr	46.013280	-121.150020	36
	POP042	SurveyorsCr	46.196480	-121.254910	35
07.05_FIFTNM	POP043	UpTroutCr	46.077410	-121.212210	51
	POP044	FifteenmileCr_wi	45.450900	-121.124360	125
07.1_DESCH	POP045	UpDeschutesR	45.261809	-121.034665	95

	POP046	DRTrouCr	44.821430	-121.087210	14
	POP047	WarmSpringsR	44.860900	-121.244380	50
	POP048	WarmSpringsRMillCr	44.865417	-121.447637	159
	POP049	WarmSpringsR_trap	44.860900	-121.244380	50
07.2_JDAY	POP053	LowJohnDayR	44.742749	-120.171219	36
	POP054	SFJohnDayR	44.286801	-119.542160	97
	POP055	NFJohnDayR	44.779154	-119.014929	41
	POP060	MFJohnDayR	44.577569	-118.805415	37
	POP063	UpJohnDayR	44.451083	-119.035553	78
07.3_UMAT	POP064	IskuulpaCr	45.699380	-118.396470	71
	POP065	UmatillaR_Minthorn	45.699380	-118.396470	74
07.4_LOWSNAKE	POP066	TouchetR	46.033994	-118.683625	58
	POP088	Alpowa Cr. '10	46.407620	-117.219784	51
	POP089	Asotin Cr. '10			59
	POP108	George Cr. '10	46.302906	-117.116818	53
	POP136	Sheep Cr. '14	45.467700	-116.555300	47
	POP143	Tucannon R. '10	46.309733	-117.657180	29
	POP144	Tucannon R. '11	46.309733	-117.657180	25
	POP145	Tucannon R. '13	46.309733	-117.657180	35
07.5_IMNA	POP068	LightningCr	45.655370	-116.726530	76
	POP109	Gumboot Cr. '12			17
	POP110	Gumboot Cr. '13			21
	POP124	Mahogany Cr. '11			6
	POP125	Mahogany Cr. '13			8
07.52_GrandeRonde	POP067	JosephCr	46.041552	-117.001710	45
	POP069	UpGrandeRondeR	45.370986	-117.865698	325
	POP070	WenahaR	45.945347	-117.451303	77
	POP112	Joseph Cr. '11	46.027769	-117.017715	55
07.6_LOWCLWR	POP091	Big Bear Cr. '12	46.630629	-116.656189	9
	POP102	E.F. Potlatch R. '16	46.798444	-116.419441	52
	POP113	Little Bear Cr. '12	46.637226	-116.677980	67
	POP123	Lapwai Cr. '13	46.334260	-116.603210	32
	POP126	Mission Cr. '13	46.317070	-116.711300	50
	POP140	Sweetwater Cr. '13	46.294630	-116.856500	65
	POP146	Webb Cr. '13	46.313980	-116.809080	30
	POP148	Potlatch R., W.F. '10	46.805376	-116.418187	50
07.7_LOWSALM	POP094	Boulder Cr. '14	45.154090	-116.395400	46
	POP137	Slate Cr. '13	45.639380	-116.120170	28
	POP149	Whitebird Cr. '14	45.770490	-116.290730	47
08_YAKIMA	POP071	AhtanumCr	46.531800	-120.681500	85
	POP072	CowicheCrowCr	46.628791	-120.570774	74
	POP073	LRattlesnakeCr	46.766026	-121.046083	70
	POP074	NFLNachesR	47.109039	-121.320081	66

09_UPPCOL	POP075	NileCr	46.861839	-121.048875	97
	POP077	QuartzCr	45.945650	-120.509653	96
	POP078	SatusCr	46.067822	-120.592854	442
	POP079.1	TaneumCr	47.092889	-120.831758	12
	POP079.2	TeanawayR	47.285885	-120.899339	17
	POP080	ToppenishCr	46.322706	-120.874878	134
	POP081	EntiatR	47.664071	-120.241689	43
	POP082	IcicleCr	47.559120	-120.674150	10
	POP083	MethowR	48.049472	-119.901200	49
	POP084	MethowH	48.476859	-120.205384	25
10_SFCLWR	POP085	NasonCr	47.801890	-120.715010	19
	POP086	OmakCr	48.395658	-119.504272	125
	POP087	WellsH	47.559120	-120.674150	11
	POP099	Clear Cr. '15			23
	POP100	Crooked R. '13	45.822237	-115.526680	56
	POP122	Lolo Cr. '12			46
	POP128	Newsome Cr. '12	45.836560	-115.615180	48
	POP141	Tenmile Cr. '13			51
	POP092	Bear Cr. '12	46.019020	-114.837580	27
	POP096	Canyon Cr. '04	46.216135	-115.555865	45
11_UPCLWR	POP097	Lochsa R. Crooked Fork '00	46.525108	-114.678649	46
	POP101	E.F. Moose Cr. '12	46.173560	-114.886080	44
	POP107	Gedney Cr. '04	46.058216	-115.313589	51
	POP114	Little Clearwater R. '08	45.753673	-114.774752	57
	POP129	Ohara Cr. '13	46.048060	-115.516940	28
	POP138	Selway R. '08			47
	POP142	Three Links Cr. '12	46.111200	-115.071900	34
	POP150	White Cap Cr. '08	45.868734	-114.728378	55
	POP103	Salmon R. S.F., E.F. '09	45.012862	-115.713109	25
	POP104	Salmon R. S.F., E.F. '10			22
12_SFSALM	POP135	Secesh R. '11			63
	POP090	Bargamin Cr. '15	45.721674	-115.034530	56
	POP093	Big Cr. '11	45.092594	-114.732175	47
	POP095	Bear Valley Cr. '11			35
	POP098	Chamberlain Cr. '16			50
	POP106	Elk Cr. '11			47
	POP131	Pistol Cr. '12	44.723520	-115.150140	38
	POP132	Rapid R. '12	44.680060	-115.152270	32
	POP147	W.F. Chamberlain Cr. '16			50
	POP105	Salmon R. E.F. '16	44.115434	-114.429996	65
14_UPSALM	POP111	Hayden Cr. '17	44.861600	-113.631880	53
	POP127	Morgan Cr. '12	44.667570	-114.230070	47
	POP130	Panther Cr. '13	45.034940	-114.299490	51

POP133	Salmon R. '18	44.163590	-114.886786	2
POP134	Salmon R. '16	44.150558	-114.885088	50
Grand Total				7422

Combined application of PBT and GSI

We combined PBT and GSI results together by first accepting all confident PBT assignments to hatchery broodstock (i.e., $\text{LOD} \geq 14$ & $\text{FDR} \leq 0.1$) (See methods for [Parentage assignments using SNPPIT software v1.0](#), ID: 1341). For the remaining individuals, we used the best estimate of GSI assignments (regardless of the probability of assignment) provided by the program ONCOR to determine likely reporting group of origin (Method: [Assigning individual samples using Individual Assignment \(IA\) genetic methods v1.0](#), ID: 1334). For the assignment of sockeye, GSI via ONCOR was used. We also have a baseline of candidate parents used in the reintroduction of Sockeye salmon which requires both SNPPIT and a program to perform single parentage assignments (SEQUOIA). For Chinook salmon, all age classes (3-, 4-, and 5+ year old fish) can be identified from Snake River and Columbia River stocks using PBT.

Use of SCOBIDEUX and SPIBETR functions to expand PBT and GSI for abundance in harvest samples

After combining PBT and GSI assignments, we examined the stratification of the harvest samples for the way in which harvest managers stratify the data to perform CREEL estimates of total harvested fish. In some cases, fisheries are mark selective and so only adipose clipped fish are necessary to sample to estimate the stock-specific abundances in the harvest. We also pooled our sample into the same temporal and spatial stratification in which harvest managers report CREEL estimates. For example, if the total harvested fish is only estimated for the entire catch without further breakdown into weeks or months, we had to treat our harvest sample as a single stratum. Ideally, fish are sampled randomly to acquire the DNA tissue samples in the same stratification units that harvest managers report. However, there are several ways in which the selection of the harvest samples has been non-random in the past. For example, the harvest monitors that work for the joint states (ODFW and WDFW) classify their samples into 3 “sample categories” in which samples can either be random without a CWT, random with a CWT, or non-random with a CWT. Because, the current management of the fisheries rely on CWT data, fish that are not part of the normal random sample for the CREEL estimates will be wanded for the presence of CWTs and if there is a positive CWT detection that fish will get sampled with biodata and genetic tissue collection. For genetic analysis, the non-random CWT fish should be excluded to obtain a random sample, but could be used for purposes of comparing CWT and PBT stock ID results. In past years, we may have inadvertently subsampled both random and non-random samples and used both types in our analyses.

We made every effort to match representative DNA sampling of these harvest management strata, which is a difference in this year’s analysis compared to previous years. Specifically, we describe our stratification for each fishery sample in the following section:

The spring test fishery:

This fishery is conducted in the early portion of the spring Chinook salmon management period and occurs in the commercial sections 2 and 3 (a portion of “Region B”). Chinook salmon are visually stock-identified (VSI) to lower river and upriver stock classifications. A fleet of boats conduct drift fishing each week and catch per unit effort (CPUE) is estimated using

the number of fish caught per number of drifts for each week. Tissue samples are obtained from both the adipose clipped and adipose intact fish caught in the test fishery at a high rate (>50%). The CPUE has appeared to be a good predictor for the timing and strength of the first peak of the run of spring chinook at Bonneville Dam (Hess et al. 2019), and in this report we established a new strategy for analysis of this sample to obtain stock-specific CPUE for both the adipose clipped and adipose intact upriver chinook salmon. For our sample, we used only the VSI-upriver chinook salmon that were caught in sections 2 and 3 of the test fishery, and stratified into pools of weekly drifts (as indicated in Table 10). Weeks were pooled to obtain sample sizes >10 fish for most strata. We applied these stratified samples to the appropriate CPUE estimated for the respective temporal strata for each of the adipose clipped and unclipped observed VIS-upriver test fish.

Table 10. The sample rate and stratification for genetic analysis of the VSI-upriver adipose clipped and unclipped adult Chinook salmon from the spring test fishery in 2018.

Week	Drifts	Observed Upriver AD			SAMPLE (Upriver VSI AD)			Observed Upriver AD-intact			SAMPLE (Upriver VSI AD-intact)		
		Pooled			Pooled			Pooled			Pooled		
		N	N	CPUE	N	N	Rate	N	N	CPUE	N	N	Rate
12	15	1	17	0.36	1	15	0.88	0	6	0.13	0	3	0.50
13	17	8			7			4			3		
14	15	8			7			2			0		
15	16	12	79	2.47	11	64	0.81	6	16	0.50	6	14	0.88
16	16	67			53			10			8		
17	16	73	121	4.32	68	116	0.96	13	26	0.93	10	11	0.42
18	12	48			48			13			1		
		217	217		195	195		48	48		28	28	

Note: Only the visual stock identified (VSI) upriver Chinook salmon were analyzed from the spring test fishery. The Catch per Unit Effort (CPUE) was calculated from the number of fish observed / the number of "Drifts" made total for the fleet of boats used in the test fishery.

The spring chinook salmon sport fishery:

This fishery is a mark-selective, which means only the adipose-clipped fish are retained ("kept" fish). All the adipose-intact fish are released, and none of these released fish are tissue sampled however a portion of them are counted as mortalities. It may be possible in the future to use the sport test fishery unclipped chinook salmon stock composition to characterize the release mortalities from the spring sport fishery because the test fishery overlaps in timing and region with the spring sport fishery. Harvest is estimated and reported for the total lower Columbia mainstem instead of being geographically stratified into regions A and B. This lack of geographic stratification could be problematic and lead to bias in the genetic analysis unless all sampling is random and occurs at the same rate across regions. We have shown in the past that there are different compositions of stocks across regions (Hess et al. 2019). Sampling conducted by the joint states is typically random and sampling rates are even across the fishing zones, which would avoid this bias.

Table 11. The sample rate and stratification for genetic analysis of the kept adipose clipped adult Chinook salmon from the spring sport fishery in 2018.

	Total Adult Chinook			Sample of Total Adult Chinook Kept			
	Kept	Rel.	Rel. Mortality	regionA	regionB	total	rate
Jan-Feb Total	18	0	0				0.12
March Total	1,871	278	28	109	121	230	
April Total	4,119	581	58	169	335	504	0.12
May Total	468	355	36	29	14	43	0.09
June 1-15 Total	1,033	316	32	40	56	96	0.09
Season Total	7,509	1,530	154	347	526	873	0.12

Note: The released fish (“Rel.”) were not sampled for genetic analysis and so only the “kept” fish could be analyzed. The samples of kept fish came from both the regions A and B but for the analysis the sample was treated as a single region for compatibility with the way the catch is reported. Monthly strata were used and the first stratum was pooled to include all fish harvested from January through March as indicated by the outlines in the table.

The spring chinook salmon platform hook and line permit fishery:

Yakama Nation executed this fishery in 2018 below Bonneville Dam. The total catch was 454 adult spring chinook salmon and samples obtained for the entire period were 273 or a sample rate of 60.1%. All samples were pooled and the catch was estimated by adipose clipped and intact stocks.

The spring chinook salmon zone 6 ceremonial permit fishery was not analyzed due to the non-representative manner in which samples were obtained. Samples are difficult to obtain in general due to the use of these fish for ceremonies. Ideally, samples would be collected from the harvest of all four member tribes (total ceremonial harvest in 2018 = 8,718) and temporal stratification would be applied; however, samples were gathered from the Yakama Nation harvest exclusively in 2018 (N=168, Table 6) and the harvested fish are not possible to report on a weekly basis or by reservoir.

The summer chinook salmon sport fishery is mark-selective similar to the spring sport fishery. The total adult fish that were kept in 2018 were 1,027 fish and of those, we obtained 72 tissue samples from adipose clipped fish across regions A and B. The summer chinook salmon are mostly destined for the upper Columbia and so stock composition should be similar across regions A and B. We pooled all 72 tissue samples which were all random (sample rate = 7%) and used the pooled sample to estimate stock composition in the fishery. There were 750 fish that were released and of those, 15% (113 fish) were assumed to be release mortalities. We did not estimate the stock composition of these release mortalities, although this could be done since the composition should be similar to the kept fish.

The summer chinook salmon Treaty fishery in zone 6 is estimated by adipose clipped and unclipped adults by statistical week (Table 12). We split our sample by adipose clip and pooled samples across weeks to obtain a minimum of 10 fish per stratum to analyze the stock composition of this harvest.

Table 12. The sample rate and stratification for genetic analysis of the adipose clipped and unclipped adult Chinook salmon from the zone 6 summer Treaty fishery in 2018.

Week	Harvest Estimate		Sample N		Sample rate	
	AD	AI	AD	AI	AD	AI
24	280	77			0.02	0.03
25	3,083	628	66	24		
26	2,228	550	41	15		
27	1,375	315	30	13	0.02	0.03
28	961	149	23	2		
29	120	34	3			
30-31	43	12	2	2		

Note: The adipose clipped (AD) and unclipped/intact (AI) fish shown by harvest estimate and sample size. Samples were pooled across weeks as indicated by the boxed samples in the table.

The fall Non-Treaty commercial fishery:

This fishery occurred in two separate weeks in August in region A and was reported by total adults and jacks. We obtained random samples of both adipose clipped and unclipped chinook salmon (both adults and jacks) and stratified by weeks to analyze the genetic stock composition of this harvest (**Table 13**). However, limited funds required subsampling. Subsampling did not keep adipose clipped and unclipped fish in the same ratios as compared to the total sample, which may have introduced some bias into the final estimates of total clipped and unclipped genetic stocks.

Table 13. The sample rate and stratification for genetic analysis of the adult and jack Chinook salmon from fall Non-Treaty commercial fishery in 2018.

Week	Harvest estimate adult+jack	Total Sample N				SubSample N			
		AD	AI	total	rate	AD	AI	total	rate
34	3351	94	100	194	0.06	57	50	107	0.03
35	4969	109	135	244	0.05	53	50	103	0.02

Note: The harvest estimate combines both adults and jacks and does not distinguish clipped and unclipped fish. The total sample was originally a random sample of the harvest but limited funds required subsampling. Subsampling did not keep adipose clipped and unclipped fish in the same ratios as compared to the total sample, which may have introduced some bias into the final estimates of total clipped and unclipped genetic stocks.

The fall Non-Treaty sport fishery:

This fishery is not mark-selective and the stock composition of the released fish is assumed to be similar to the kept fish. We stratified the samples of kept fish by month and estimated the stock composition of the adipose clipped and adipose intact fish using the sample data (Table 14). The catch estimate for this fishery is reported by month for the adult fish and further is not stratified geographically into regions. We recorded geographic region in the sample data and so this geographic stratification could be applied if the harvest estimates were stratified similarly. This geographic stratification should only be necessary if the sample were not obtained randomly with respect to time and region. However, we used only the random samples obtained from this fishery.

Table 14. The sample rate and stratification for genetic analysis of the kept adult Chinook salmon from the fall Non-Treaty sport fishery in 2018.

Catch Estimate				Sample of kept		
Month	kept Adult	Released	Release mortality	AD	AI	rate
Aug	4119	232	35	26	72	0.02
Sep	5683	645	97	37	115	0.03
	9802			63	187	0.03

Note: The sample numbers of the kept Chinook salmon were stratified by month and adipose clip data was used to estimate the stock composition of both adipose clipped (AD) and adiposed intact (AI) stocks.

The fall Treaty commercial fishery:

This fishery occurred in zone 6 across several weeks of the fall management period. This fishery is monitored by visually identifying tules versus bright stocks and estimating each separately (Table 15). Tissue collection is mostly obtained from the fish identified as bright stocks, and very few tule stocks are sampled. Therefore, genetic analysis was constrained to estimating stock composition within the visually-identified bright harvest.

Table 15. The sample rate and stratification for genetic analysis of the adult bright Chinook salmon from fall Treaty zone 6 commercial fishery in 2018.

Week	Harvest estimate		Sample of Brights		
	Brights	Tules	AD	AI	rate
31 - 34	652	19	3	9	0.01
35	5,365	3319	25	53	
36	11,530	6110	49	51	0.01
37	10,717	3102	65	100	0.02
38	5,941	1206	32	89	0.02
39	3,962	940	39	98	0.03
40	2,786	493	15	54	0.02
total	40,953	15,189	228	454	0.02

The fall test fishery or pound net fishery:

The pound net is a relatively recent gear type that is being developed by the joint states. In 2018, the fall pound net fishery was conducted in the Cathlamet Channel (area 83) and the numbers and genetic samples of the kept (Table 16) and released (Table 17) Chinook salmon were stratified to obtain at least 10 fish in each stratum. The Chinook jacks were not expanded for stock composition due to the low sample size.

Table 16. The sample rate and stratification for genetic analysis of the kept adipose clipped Chinook salmon from the fall pound net fishery in 2018.

2018 Pound Net Trap Kept Catch Summary (all adipose-clipped)

Week	Sample			Sample		
	Chin Adults	Chin Adults	Rate	Chin Jacks	Chin Jacks	Rate
34	47	1	0.46	2		0.21
35	353	182		28	8	
36	105	52	0.50	5	1	
37	92	52	0.57	13	1	
38	3	2	0.67	0		
	600	289	0.48	48	10	0.21

Note: The “Sample” indicates numbers of successfully genotyped fish in each stratum. The weekly strata were pooled as indicated by the outlined boxes to obtain at least 10 fish in each stratum. The “Rate” is the sample rate attained for each pooled stratum.

Table 17. The sample rate and stratification for genetic analysis of the released adipose unclipped adult Chinook salmon from the fall pound net fishery in 2018

2018 Pound Net Trap Released Catch Summary (adipose unclipped)

Week	Chin Adults	Sample	
	No Ad-clip		
34	70	21	0.30
35	445	88	0.20
36	152	28	0.18
37	223	33	0.15
38	11	2	
41	0		
42	1		
43	1		
	903	172	0.19

Note: The “Sample” indicates numbers of successfully genotyped fish in each stratum. The weekly strata were pooled as indicated by the outlined boxes to obtain at least 10 fish in each stratum. The “Rate” is the sample rate attained for each pooled stratum.

The Sockeye Non-Treaty sport fishery:

This fishery was estimated to have 73 and 212 kept and released fish, respectively, for a total mortality of 111 sockeye. We successfully genotyped a total of 9 of the kept fish which were assumed representative of the total mortality (sample rate = 8.1% of the total mortality). All samples were pooled to estimate stock composition of this harvest. Every fish in the sample assigned to the Osoyoos stock (i.e., 100% of the 111 sockeye mortalities were estimated to be Osoyoos).

The Sockeye Treaty commercial fishery:

This fishery occurred in zone 6 and catch estimates were reported by week in the summer and additional estimates of sockeye were reported in zone 6 during the spring and fall periods as well as below Bonneville Dam. All Sockeye salmon catch estimates were combined for the total harvest (**Table 18**).

Table 18. The sample rate and stratification for genetic analysis of the harvested Sockeye salmon from the Treaty fishery in 2018.

Week	Harvest	Sample							
	estimate	LBC	Odell	Osoyoos	Redfish	Wenatchee	Yakima	total	rate
21-24	299			4				4	0.02
25	2,346			40		11		51	
26	2,494			33	3	13	1	50	0.02
27	1,546		1	40		4		45	0.03
28	837			32		2		34	0.04
29	121	1		25		3	2	31	0.19
30	83			7		1		8	
31	0								
total	7,727	1	1	181	3	34	3	223	0.03

Results

Information content of the 254 SNPs for Chinook salmon PBT applications

The 93 SNPs that have been used since the beginning of PBT applications in the Columbia River basin were originally optimized for Spring Chinook Salmon broodstocks. One issue that has recently become problematic (Hess et al. 2019) is that there are individual genotypes that have been causing the software SNPPIT to crash. We rely on this software for our PBT analyses and so resolving this problem is critical to the effectiveness of PBT applications in the long term. We examined a set of 299 SNPs in the GT-seq panel for information content (observed heterozygosity) in each of the three major lineages of Chinook Salmon represented by a set of broodstocks collected in 2015 (Hess et al. 2019). We first discovered that among the original 93 SNPs, there were 92 that were found above an observed heterozygosity of 10% for the stream-type lineage broodstocks of spawn year 2015 (Table 12 in Hess et al. 2019). In contrast, there were 72 and 64 SNPs with observed heterozygosities >10% for the ocean-type and Lower Columbia lineages, respectively. According to Steele et al. (2013), less than 72 highly polymorphic SNPs is the point at which accuracy for PBT was observed to decline. We suspect that it is the lack of sufficient numbers of SNP markers with high information content (i.e. observed heterozygosity) that has been causing SNPPIT to crash when we analyze parent-offspring pairs from ocean-type and Lower Columbia River broodstocks with low polymorphism at these 93 SNPs.

To resolve this issue, we recommend the use of a larger set of SNP markers (~200) that have greater than 10% observed heterozygosity in each of the major lineages of Chinook Salmon. Our evaluation of the broodstocks from SY2015 indicated a set of 269 SNP loci that had >10% observed heterozygosity for at least 1 major lineage, and within this panel of 269 loci there were 193, 218, and 195 SNPs with >10% observed heterozygosity for each of the major lineages stream-type, ocean-type, and Lower Columbia, respectively (Hess et al. 2019). Based on this analysis, a set of 254 SNPs were chosen from these 269 loci and overlap with our current GT-seq panel (Table 19).

Application of the 254 SNP panel to the Chinook Salmon passing Bonneville Dam in 2018 revealed that the use of this expanded panel fully resolved the crashing issue with SNPPIT.

904 This result confirmed that the source of the crashing issue was likely related to the lack of
905 sufficient numbers of SNPs with high information content that is primarily affecting ocean-type
906 broodstocks. Further, the higher number of SNP loci requires decreased computational time to
907 run the parentage analyses, which is extremely helpful for the relatively short time available to
908 process in-season samples.

909 **Table 19. The “legacy set” of 93 SNP loci and the set of 254 SNP loci with high information**
910 **content used for PBT applications of the three major genetic lineages of Columbia River**
911 **Chinook salmon.**

SNP93 order	Best 254	Locus
1	1	Ots_100884287
2	2	Ots_101554407
3	3	Ots_101704143
4	4	Ots_102414395
5	5	Ots_102801308
6	6	Ots_103122180
7	7	Ots_10441588
8	8	Ots_105105613
9	9	Ots_105132200
10	10	Ots_105385421
11	11	Ots_105407117
12	12	Ots_108820336
13	13	Ots_109525816
14	14	Ots_110064383
15	15	Ots_110201363
16	16	Ots_110495380
17	17	Ots_11055164
18	18	Ots_110689218
19	19	Ots_11230143
20	20	Ots_112419131
21	21	Ots_112820284
22	22	Ots_112876371
23	23	Ots_113242216
24	24	Ots_115987325
25	25	Ots_117432409
26	26	Ots_11820561
27	27	Ots_118938325
28	28	Ots_123921111
29	29	Ots_124774477
30	30	Ots_12875761R
31	31	Ots_129458451
32	32	Ots_94857232R
33	33	Ots_9490399R
34	34	Ots_96500180
35	35	Ots_96899357R
36	36	Ots_ARNT
37	37	Ots_AsnRS60
38	38	Ots_brp1664

39	39	Ots_CD592
40	40	Ots_CirpA
41	41	Ots_cox1241
42	42	Ots_E2275
43	43	Ots_Est740
44	44	Ots_ETIF1A
45	45	Ots_FGF6B_1
46	46	Ots_GCSH
47	47	Ots_GDH81x
48	48	Ots_GPH318
49	49	Ots_GTH2B550
50	50	Ots_HMGB173
51	51	Ots_hsc713488
52	52	Ots_HSP90B100
53	53	Ots_IGFI176
54	54	Ots_Ikaros250
55	55	Ots_IL8R_C8
56	56	Ots_mapK3309
57	57	Ots_mapKpr151
58	58	Ots_MHC2
59	59	Ots_mybp85
60	60	Ots_NFYB147
61	61	Ots_nkef192
62	62	Ots_NOD1
63	63	Ots_ntl255
64	64	Ots_OTALDBINT1SNP1
65	65	Ots_OTDESMIN19SNP1
66	66	Ots_OTSTF1SNP1
67	67	Ots_P53
68	68	Ots_parp3286
69	70	Ots_pop596
70	71	Ots_ppie245
71	72	Ots_Prl2
72	73	Ots_RAG3
73	74	Ots_redd1187
74	75	Ots_S71
75	76	Ots_SClkF2R2135
76	77	Ots_SWS1op182
77	78	Ots_TAPBP
78	79	Ots_TGFB
79	80	Ots_Thio
80	81	Ots_TLR3
81	82	Ots_tpx2125

82	83	Ots_txnip321
83	84	Ots_u0707161
84	85	Ots_u0717135
85	86	Ots_u0718378
86	87	Ots_u0725325
87	88	Ots_u0749290
88	89	Ots_u100275
89	90	Ots_u21185
90	91	Ots_u492
91	92	Ots_u675
92	93	Ots_unk526
93	94	Ots_vatf251
	69	Ots_pigh105
	95	Ots_106747239
	96	Ots_11345740R
	97	Ots_123048521
	98	Ots_96222525
	99	Ots_97077179R
	100	Ots_AldB1122
	101	Ots_arp436
	102	Ots_aspat196
	103	Ots_Cath_D141
	104	Ots_CD63
	105	Ots_CRB211
	106	Ots_EndoRB1486
	107	Ots_EP529
	108	Ots_FARSLA220
	109	Ots_GST207
	110	Ots_GST375
	111	Ots_hsp27b150
	112	Ots_IL11
	113	Ots_myo1a384
	114	Ots_myoD364
	115	Ots_nramp321
	116	Ots_Ots311101x
	117	Ots_P450
	118	Ots_PGK54
	119	Ots_RFC2558
	120	Ots_SL
	121	Ots_u0720332
	122	Ots_u0753133
	123	Ots_u0757120
	124	Ots_u202161

125	Ots_ZR575
126	Ots_LWSop638
127	Ots_u1008108
128	Ots_IsoT
129	Ots_101119381
130	Ots_102213210
131	Ots_102457132
132	Ots_102867609
133	Ots_10649970
134	Ots_107074284
135	Ots_10728593
136	Ots_107806821
137	Ots_108007208
138	Ots_108390329
139	Ots_108735302
140	Ots_109693392
141	Ots_111681657
142	Ots_112208722
143	Ots_117242136
144	Ots_117259271
145	Ots_118175479
146	Ots_12241456
147	Ots_12723662
148	Ots_12830257
149	Ots_128693461
150	Ots_129144472
151	Ots_13072099
152	Ots_131460584
153	Ots_131906141
154	Ots_99550204
155	Ots_DDX5171
156	Ots_Est1363
157	Ots_HFABP34
158	Ots_hnRNPL533
159	Ots_Hsp90a
160	Ots_nelfd163
161	Ots_OTSMTASNP1
162	Ots_P450288
163	Ots_stk6516
164	Ots_TCTA58
165	Ots_u1007124
166	Ots_U2362227
167	Ots_U2362330

168	Ots_U2446123
169	Ots_unk110438
170	Ots_unk183239
171	Ots_unk351349
172	Ots_unk793650
173	Ots_unk948051
175	Ots_104063132
176	Ots_105401325
177	Ots_106313729
178	Ots_106419b618
179	Ots_107607315
180	Ots_110381164
181	Ots_117370471
182	Ots_120950417
183	Ots_126619400
184	Ots_127760569
185	Ots_12987055
186	Ots_131802393
187	Ots_95442b204
188	Ots_9766056
189	Ots_afmid196
190	Ots_AldoB4183
191	Ots_BMP2SNP1
192	Ots_cgo2422
193	Ots_crRAD1044725
194	Ots_crRAD1162055
195	Ots_crRAD1203739
196	Ots_crRAD1372551
197	Ots_crRAD1654050
198	Ots_crRAD1752758
199	Ots_crRAD1849265
200	Ots_crRAD1893760
201	Ots_crRAD2026246
202	Ots_crRAD2037666
203	Ots_crRAD2088770
204	Ots_crRAD2111524
205	Ots_crRAD2296032
206	Ots_crRAD2480774
207	Ots_crRAD2536750
208	Ots_crRAD25559
209	Ots_crRAD2616569
211	Ots_crRAD2716455
212	Ots_crRAD2751569

213	Ots_crRAD280642
214	Ots_crRAD3349171
215	Ots_crRAD3439733
216	Ots_crRAD3531366
217	Ots_crRAD3607229
218	Ots_crRAD3615244
219	Ots_crRAD4458867
220	Ots_crRAD4608156
221	Ots_crRAD4675142
222	Ots_crRAD4729755
223	Ots_crRAD5547526
224	Ots_crRAD5752066
225	Ots_crRAD5768734
226	Ots_crRAD6061446
227	Ots_crRAD6062051
228	Ots_crRAD6152371
229	Ots_crRAD6633060
230	Ots_crRAD6932753
231	Ots_crRAD7382360
232	Ots_crRAD7476628
233	Ots_crRAD7558170
234	Ots_crRAD7651228
235	Ots_crRAD7896846
236	Ots_crRAD9242025
237	Ots_crRAD961569
238	Ots_MetA
239	Ots_NAML12SNP1
240	Ots_Ostm1
241	Ots_PEMT
242	Ots_RAD454352
243	Ots_sept978
244	Ots_slc7a271
245	Ots_trnau1ap86
247	Ots_U212158
248	Ots_U230563
249	Ots_U2567104
250	Ots_U5049250
251	Ots_U512134
252	Ots_crRAD1828933
253	Ots_crRAD5540059
254	Ots_crRAD5737668
255	Ots_10304152
256	Ots_111084b619

257 Ots_129170683

912 Note: The loci are sorted by their order in the original panel of 93SNPs used for PBT applications, and “Best 254”
913 indicates the group of 254 SNPs that were observed with a heterozygosity >10% for at least one of the major genetic
914 lineages based on hatchery broodstock of spawn year 2015 (Hess et al. 2019).

Comparison of Coded-wire tags and PBT assignments

There were 101 coded-wire tags (CWTs) recovered and identified to hatchery stock and broodyear (BY) among the snouts recovered from the lower river fisheries (Table 20), and 77 of these CWTs also were PBT assigned (76%). Of the 77 fish with both CWT and PBT, there were 65 fish (84%) that appeared concordant with the PBT assignments according to both the hatchery source and the broodyear. There were 12 of the 77 fish with both CWT and PBT that were discordant for either hatchery source (N=6) or broodyear (N=5), and a single fish that was discordant for both hatchery source and broodyear information.

Table 20. Comparison of coded-wire tags with parentage-based tags that were identified in the lower river fisheries in 2018.

Period	Coded-wire tag		Parentage-based tagging assignment				HatcheryCorrect	BYCorrect	Missing
	Hatchery	BY	Code	Hatchery	BY	N			
01Spring	CHELAN FALLS HATCHERY	2012	OtsEASTBK12_su	Eastbank Fish Hatchery	2012	1	Y	Y	0
01Spring	CHELAN FALLS HATCHERY	2014	OtsEASTBK14_su	Eastbank Fish Hatchery	2014	4	Y	Y	0
01Spring	CHIEF JOSEPH HATCHERY	2013	OtsCJH13int_su	Chief Joseph Hatchery	2013	1	Y	Y	0
01Spring	CLE ELUM HATCHERY	2014	OtsYR14int_sp	Yakima River Roza Dam	2014	1	Y	Y	0
01Spring	DRYDEN POND	2014	OtsEASTBK14_su	Eastbank Fish Hatchery	2014	1	Y	Y	0
01Spring	LOOKINGGLASS HATCH	2014	OtsLSTW14S_sp	Lookingglass Fish Hatchery	2014	1	Y	Y	0
01Spring	LOOKINGGLASS HATCH	2015	OtsLSTW15S_sp	Lookingglass Fish Hatchery	2015	1	Y	Y	0
01Spring	MCCALL HATCHERY	2014	OtsMCCA14S_spsu	McCall Fish Hatchery	2014	2	Y	Y	0
01Spring	NASON CREEK ACC FACILITY	2015	OtsEASTBK15_sp	Eastbank Fish Hatchery	2015	1	Y	Y	0
01Spring	ROUND BUTTE HATCHERY	2014	OtsPFF14_sp	Parkdale Fish Facility	2014	2	Y	Y	0
01Spring	SIMILKAMEEN HATCHERY	2012	OtsEASTBK12_su	Eastbank Fish Hatchery	2012	1	Y	Y	0
01Spring	SIMILKAMEEN HATCHERY	2014	OtsCJH14int_su	Chief Joseph Hatchery	2014	4	Y	Y	0
01Spring	WELLS HATCHERY	2013	OtsWELLS13_su	Wells Fish Hatchery	2013	1	Y	Y	0
01Spring	WELLS HATCHERY	2014	OtsWELLS14_su	Wells Fish Hatchery	2014	12	Y	Y	0

02Summer	Carlton Acclimaiton Pond (Methow R)	2014	OtsEASTBK14_su	Eastbank Fish Hatchery	2014	1	Y	Y	0
02Summer	Carlton Acclimaiton Pond (Methow R)	2013	OtsEASTBK13_su	Eastbank Fish Hatchery	2013	1	Y	Y	0
02Summer	CHELAN FALLS HATCHERY	2012	OtsEASTBK12_su	Eastbank Fish Hatchery	2012	1	Y	Y	0
02Summer	CHELAN FALLS HATCHERY	2014	OtsEASTBK14_su	Eastbank Fish Hatchery	2014	4	Y	Y	0
02Summer	CHELAN FALLS HATCHERY	2015	OtsEASTBK15_su	Eastbank Fish Hatchery	2015	1	Y	Y	0
02Summer	CHIEF JOSEPH HATCHERY	2013	OtsCJH13seg_su	Chief Joseph Hatchery	2013	1	Y	Y	0
02Summer	CHIEF JOSEPH HATCHERY	2014	OtsCJH14seg_su	Chief Joseph Hatchery	2014	1	Y	Y	0
02Summer	DRYDEN POND	2013	OtsEASTBK13_su	Eastbank Fish Hatchery	2013	1	Y	Y	0
02Summer	DRYDEN POND	2014	OtsEASTBK14_su	Eastbank Fish Hatchery	2014	2	Y	Y	0
02Summer	ENTIAT NFH	2014	OtsENFH14_su	Entiat National Fish Hatchery	2014	2	Y	Y	0
02Summer	MCCALL HATCHERY	2014	OtsMCCA14S_spsu	McCall Fish Hatchery	2014	1	Y	Y	0
02Summer	SIMILKAMEEN HATCHERY	2014	OtsCJH14int_su	Chief Joseph Hatchery	2014	2	Y	Y	0
02Summer	WELLS HATCHERY	2013	OtsWELLS13_su	Wells Fish Hatchery	2013	1	Y	Y	0
02Summer	WELLS HATCHERY	2014	OtsWELLS14_su	Wells Fish Hatchery	2014	1	Y	Y	0
03Fall	Bonneville Hatchery	2014	OtsLWS14_fa	Little White Salmon National Fish Hatchery	2014	2	Y	Y	0
03Fall	CHIEF JOSEPH HATCHERY	2014	OtsCJH14seg_su	Chief Joseph Hatchery	2014	1	Y	Y	0
03Fall	Irrigon Hatchery	2016	OtsLYON16S_1_fa	Lyons Ferry Fish Hatchery	2016	1	Y	Y	0
03Fall	Klickitat Hatchery	2015	OtsLWS15_fa	Little White Salmon National Fish Hatchery	2015	1	Y	Y	0
03Fall	Lyons Ferry Hatchery	2014	OtsLYON14S_1_fa	Lyons Ferry Fish Hatchery	2014	4	Y	Y	0
03Fall	Priest Rapids Hatchery	2015	OtsPRH15_fa	Priest Rapids Hatchery	2015	1	Y	Y	0
03Fall	Spring Creek Hatchery	2015	OtsSPCR15_fa	Spring Creek National Fish Hatchery	2015	1	Y	Y	0
03Fall	Umatilla Hatchery	2015	OtsLWS15_fa	Little White Salmon National Fish Hatchery	2015	1	Y	Y	0
01Spring	WELLS HATCHERY	2013	OtsWELLS14_su	Wells Fish Hatchery	2014	1	Y	N	0
01Spring	WELLS HATCHERY	2014	OtsWELLS13_su	Wells Fish Hatchery	2013	1	Y	N	0
02Summer	Carlton Acclimaiton Pond (Methow R)	2013	OtsCJH14int_su	Chief Joseph Hatchery	2014	1	Y	N	0
02Summer	DRYDEN POND	2013	OtsEASTBK12_su	Eastbank Fish Hatchery	2012	1	Y	N	0
02Summer	SIMILKAMEEN HATCHERY	2014	OtsEASTBK13_su	Eastbank Fish Hatchery	2013	1	Y	N	0
01Spring	CHELAN FALLS HATCHERY	2014	OtsCJH14int_su	Chief Joseph Hatchery	2014	1	N	Y	0
01Spring	LOOKINGGLASS HATCH	2014	OtsPFF14_sp	Parkdale Fish Facility	2014	1	N	Y	0

02Summer	Sandy Hatchery	2014	OtsEASTBK14_su	Eastbank Fish Hatchery	2014	2	N	Y	0
02Summer	Sandy Hatchery	2014	OtsENFH14_su	Entiat National Fish Hatchery	2014	1	N	Y	0
03Fall	Bonneville Hatchery	2014	OtsLYON14S_1_fa	Lyons Ferry Fish Hatchery	2014	1	N	Y	0
02Summer	MCKENZIE HATCHERY	2013	OtsENFH14_su	Entiat National Fish Hatchery	2014	1	N	N	0
01Spring	DRYDEN POND	2013	NA	NA	NA	1			1
01Spring	ENTIAT NFH	2014	NA	NA	NA	1			1
01Spring	LEWIS RIVER HATCHERY	2015	NA	NA	NA	1			1
01Spring	LOOKINGGLASS HATCH	2014	NA	NA	NA	1			1
01Spring	MCKENZIE HATCHERY	2014	NA	NA	NA	1			1
02Summer	CHELAN FALLS HATCHERY	2013	NA	NA	NA	1			1
02Summer	Colitz Salmon Hatchery	2014	NA	NA	NA	1			1
02Summer	DRYDEN POND	2013	NA	NA	NA	1			1
02Summer	MCKENZIE HATCHERY	2013	NA	NA	NA	1			1
02Summer	Minto Ponds (N Santiam R)	2013	NA	NA	NA	1			1
03Fall	Klickitat Hatchery	2013	NA	NA	NA	1			1
03Fall	Sandy Hatchery	2016	NA	NA	NA	1			1
01Spring	Sandy Hatchery	2014	NA	NA	NA	2			2
01Spring	CHELAN FALLS HATCHERY	2013	NA	NA	NA	4			4
02Summer	Sandy Hatchery	2014	NA	NA	NA	6			6

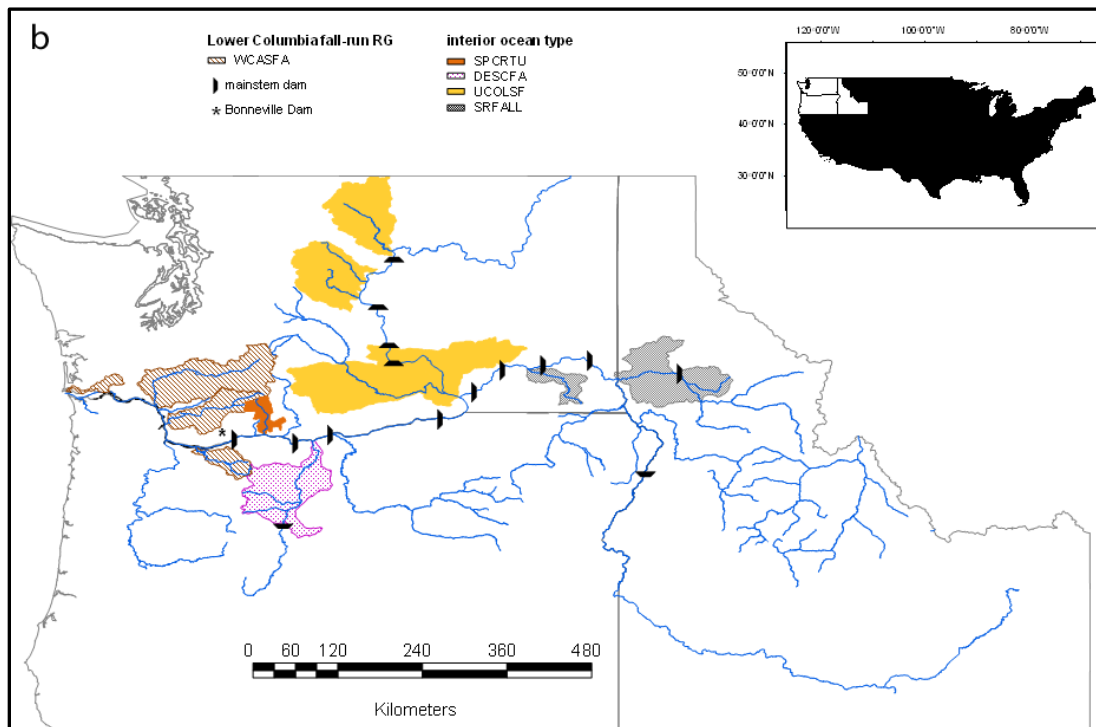
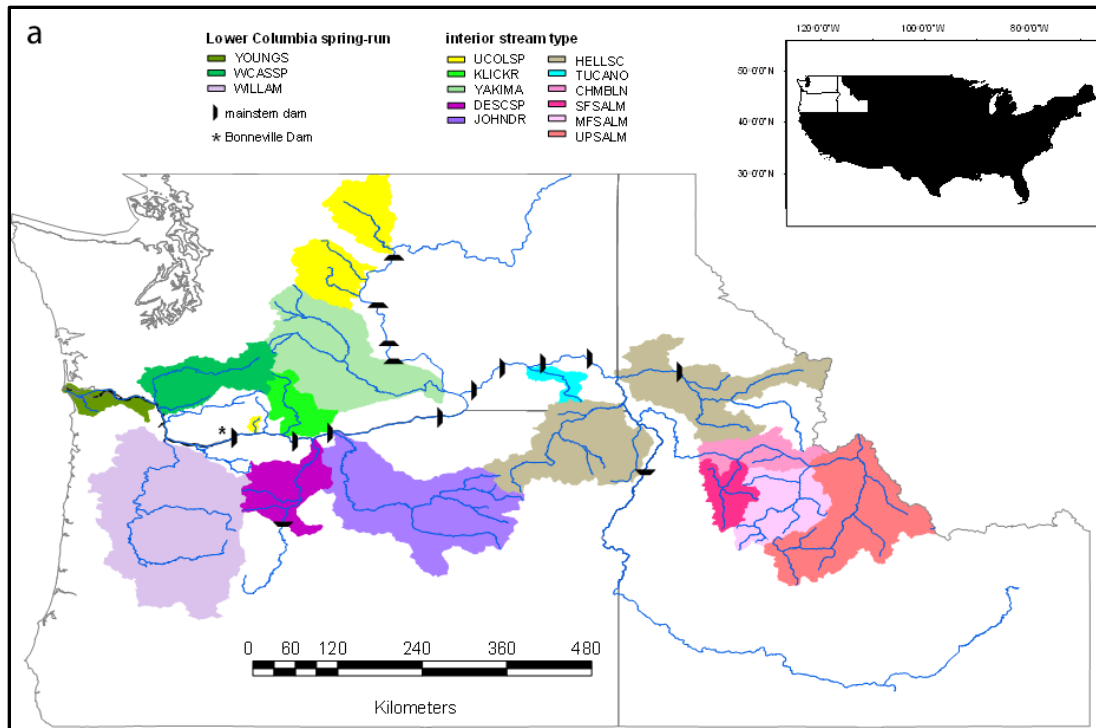
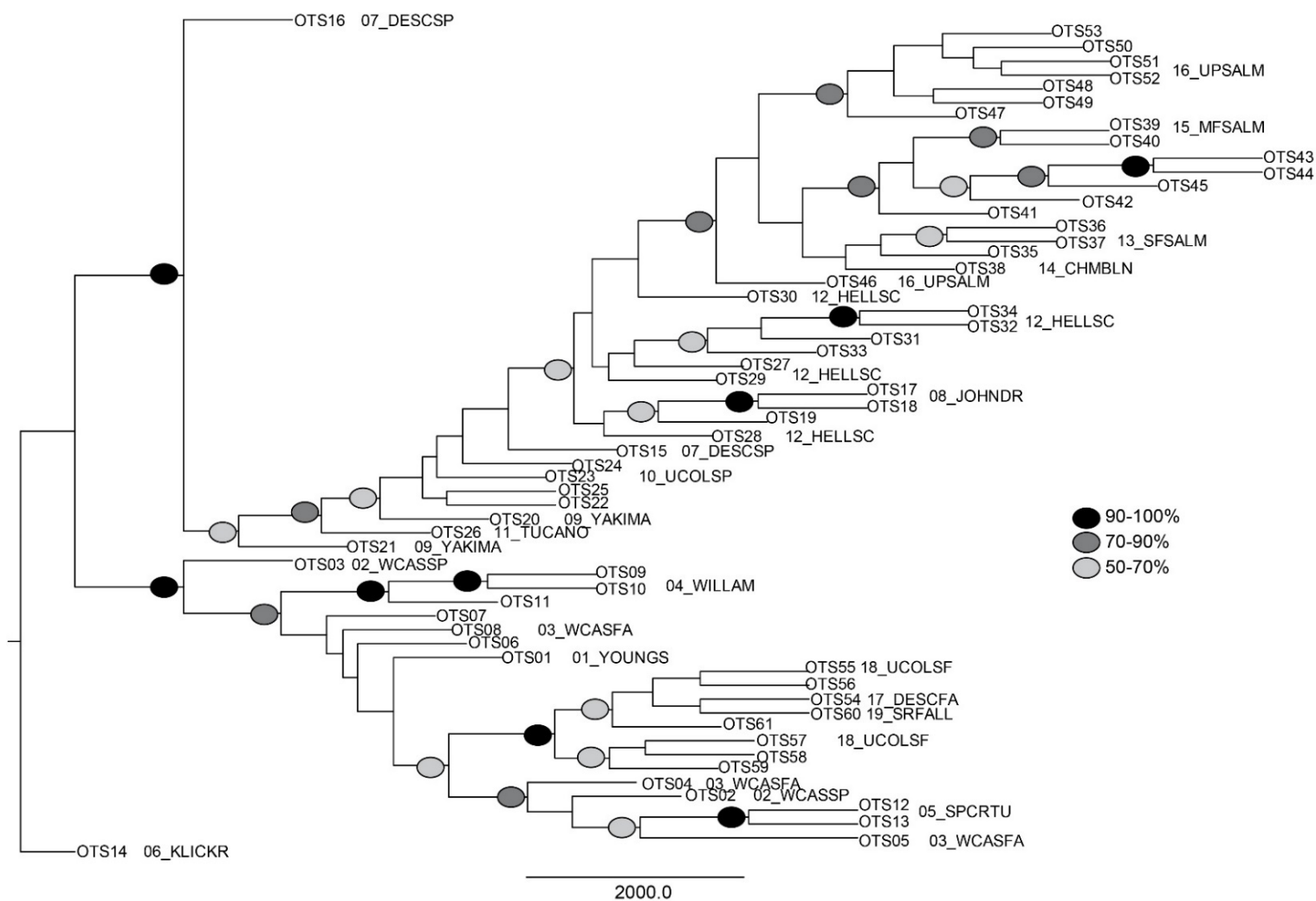


Figure 14. Map of Chinook salmon GSI reporting groups for a) Lower Columbia (LC) and interior stream type (ST) lineage, and b) interior ocean type (OT) lineage.



930 **Figure 15. Neighbor-joining tree of Chinook salmon baseline populations using Nei's 1972 genetic distance of 179 SNP loci.**

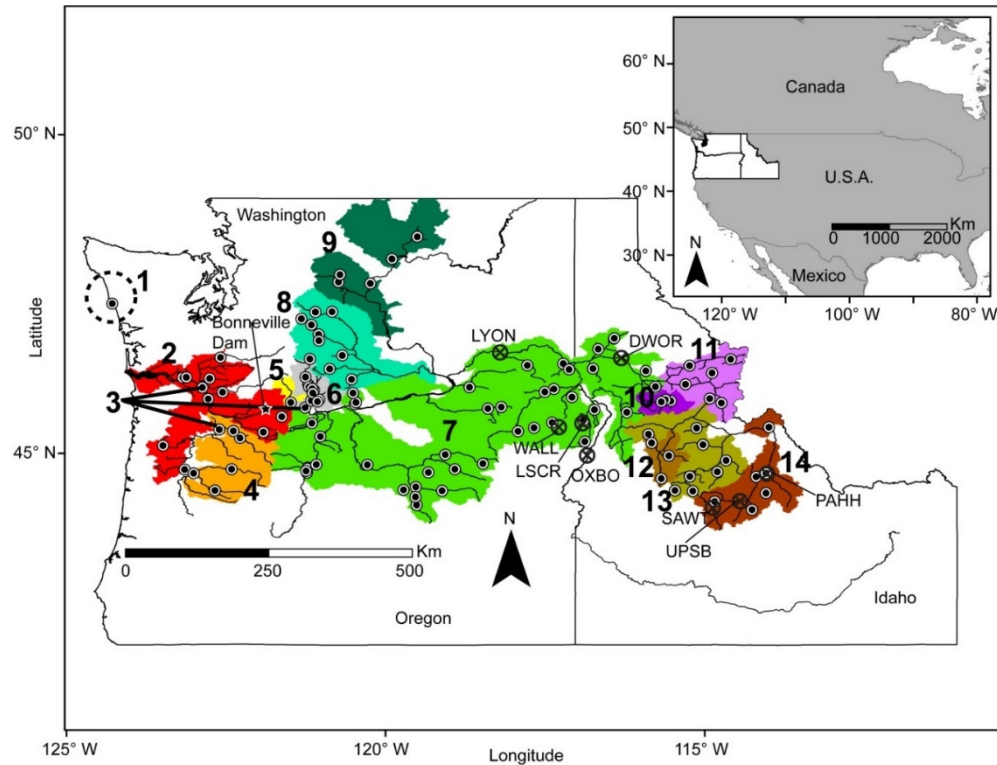


Figure 16. Geographic distribution of collections represented in the Columbia River steelhead GSI and PBT genetic baselines.

In Figure 16, the shape overlay represents the geographic extent of the following 14 reporting groups in the GSI baseline: 1) Quinault (WCOAST), 2) lower Columbia River (LOWCOL), 3) Skamania hatchery releases at three sites in lower Columbia River, Willamette River, and Klickitat River (SKAMAN), 4) Willamette River (WILLAM), 5) Big White Salmon River (BWSALM), 6) Klickitat River (KLICKR), 7) middle Columbia River, Grande Ronde River, Imnaha River, lower Snake River, lower Clearwater River, and lower Salmon River (MGILCS), 8) Yakima River (YAKIMA), 9) upper Columbia River (UPPCOL), 10) South Fork Clearwater River (SFCLWR), 11) upper Clearwater River (UPCLWR), 12) South Fork Salmon River (SFSALM), 13) Middle Fork Salmon River (MFSALM), and 14) upper Salmon River (UPSALM). There are 116 collections (filled circles) categorized into reporting groups. The PBT baseline is indicated as 8 stocks (crossed circles) corresponding to the following sites where fish are collected and spawned for broodstock: Lyons Ferry Hatchery (LYON), Wallowa (WALL), Little Sheep Creek (LSCR), Oxbow Hatchery (OXBO), Dworshak Hatchery (DWOR), upper Salmon River B-run (UPSB), Sawtooth Hatchery (SAWT), and Pahsimeroi Hatchery (PAHH). Bonneville Dam (star) is the site where fish were non-lethally sampled for the mixed-stock analysis.

Steelhead 177 SNP and 335 SNP baselines

For steelhead, we have used GSI baseline v3.3 that comprises 116 collections from throughout the Columbia River basin that are partitioned into 14 reporting groups (N= 9991) (Figure 16). However, a set of 335 SNP loci is now available for a similar set of collections representing 13 of the 14 reporting groups (N=7422 individuals, **Table 9**). We tested the accuracy of this new panel of 335 SNP loci by performing leave-1-out tests. Specifically, we generated the following different subsets of loci (Figure 17): 180 SNPs from baseline v3.3 (“Sub180”), 339 SNPs that include putatively neutral and adaptive loci (“Sub339”), 335 SNPs that exclude the run timing candidate SNPs (“Sub335NRT”), 301 SNPs that were putatively neutral (“Sub301Neu”), and 55 SNPs that were putatively adaptive (“Sub55Adapt”). We found that the 335 and 339 SNPs both performed best, but we felt it would be better to exclude the run timing candidate SNPs to avoid any unintended bias for stocks containing within population variation in run timing. Therefore, we propose testing of the 335 SNP baseline to determine whether it can replace the original 177 SNP v3.3. baseline.

However, recent testing using comparisons of PBT and GSI assignments have highlighted some potential accuracy issues with the new 335 SNP baseline. First, this new baseline is not able to accurately assign smaller subgroups within the MCGILCS reporting group (Table 8), which is one of the primary reasons it was developed. Second, the original set of reporting groups may not have improved with this new panel of SNPs. The PBT broodstocks can be categorized into “expected” reporting groups according to where they are located and which genetic stock they use for broodstock. We found that the original 177 SNP baseline was more accurate for one stock in particular, Upper Salmon River, and assigned a greater number of PBT assignments (82%) to this Upper Salmon River stock compared to the new SNP baseline of 335 SNPs (35% assigned correctly, **Table 21**, **Table 22**). Although assignment accuracies to the other reporting groups we could analyze in this way were similar between SNP panels (**Table 22**), the poor accuracy of the Upper Salmon River stock prevents us from using this baseline to replace the version 3.3 baseline until more testing can be conducted. Therefore, we continue to use the panel of 177SNPs (version 3.3) for the GSI applications in this report.

We detected samples (n=28, 1.5%) from the coastal lineage (i.e., 01_WCOAST – 06_KLICKR reporting groups) that were incorrectly assigned to reporting groups of the inland lineage (i.e., 07_MGILCS – 14_UPSALM reporting groups (**Table 20**Figure 17). We similarly detected samples (n=29, 0.34%) from reporting groups from the inland lineage were incorrectly assigned to coastal lineage reporting groups (**Table 20**). Incorrect assignments were typically distributed across multiple reporting groups, but the largest proportion of incorrect assignments were consistently to three reporting groups (07_MGILCS, 06_KLICKR, and 05_BWSALM). Samples from the geographically large 07_MGILCS reporting group were incorrectly assigned to every reporting group except 01_WCOAST, 03_SKAMAN, and 04_WILLAM (**Table 20**).

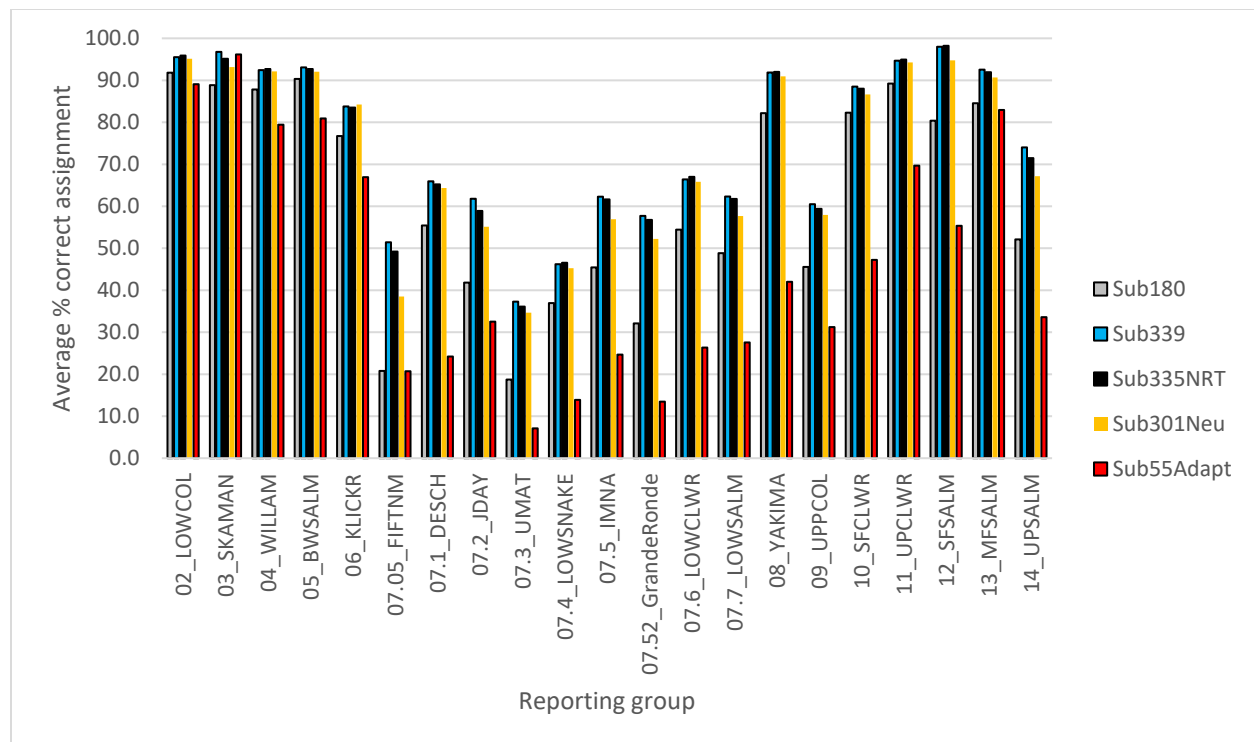


Figure 17. Proportion of steelhead in leave-one-out tests that assigned correctly for each reporting group by lineage using the following subsets of SNPs: 180 SNPs from baseline v3.3 (“Sub180”), 339 SNPs that include putatively neutral and adaptive loci (“Sub339”), 335 SNPs that exclude the run timing candidate SNPs (“Sub335NRT”), 301 SNPs that were putatively neutral (“Sub301Neu”), and 55 SNPs that were putatively adaptive (“Sub55Adapt”).

996 **Table 21. Comparison of PBT expected reporting groups versus the observed reporting groups using 177 SNPs (baseline 3.3)**
 997 **based on the assignments from the Bonneville 2019 mixture.**

PBT Expected GSI	Observed GSI						Total	Correct%
	03_SKAMAN	07_MGILCS	09_UPPCOL	10_SFCLWR	11_UPCLWR	14_UPSALM		
03_SKAMAN	22						22	100.0%
07_MGILCS		100	2			13	115	87.0%
09_UPPCOL		5	4			1	10	40.0%
10_SFCLWR		7		209	1	3	220	95.0%
14_UPSALM		17	1	1		88	107	82.2%

998
 999 **Table 22. Comparison of PBT expected reporting groups versus the observed reporting groups using 335 SNPs (new baseline)**
 1000 **based on the assignments from the Bonneville 2019 mixture.**

PBT Expected GSI	Observed GSI						Total	Correct%
	03_SKAMAN	07_MGILCS	09_UPPCOL	10_SFCLWR	11_UPCLWR	14_UPSALM		
03_SKAMAN	22						22	100.0%
07_MGILCS		112	2	1			115	97.4%
09_UPPCOL		3	7				10	70.0%
10_SFCLWR		3		216	1		220	98.2%
14_UPSALM		68	2			37	107	34.6%

1003 *Sockeye (363 SNPs) parent baseline for identifying reintroduced stocks from Yakima River*

1004 This is the first year we have been able to utilize candidate parents that were genotyped from tissues collected from carcass
 1005 spawning surveys and directly from the fish translocated from Priest Rapids Dam and released into the Yakima River (**Table 23**). A
 1006 combination of parent-pair assignments (trio assignment) and single parent assignments were performed to obtain as large of a sample
 1007 of offspring as possible. Tag rates assumed the ability to perform single parent assignments.

1008

1009

1010 **Table 23. The numbers of candidate parents genotyped relative to the number of returning and translocated adult sockeye to**
 1011 **the Yakima River each year.**

Year	Outplants	Prosser	Roza	O+P	O+R	Genotypes		Max YR	Tag rate	
						383	88		383	88
2009	1000	12	17	1012	1017	0	0	1017	0.0%	0.0%
2010	2500	11	40	2511	2540	0	0	2540	0.0%	0.0%
2011	4000	0	13	4000	4013	0	29	4013	0.0%	1.4%
2012	10000	47	154	10047	10154	0	985	10154	0.0%	18.5%
2013	4500	696	691	5196	5191	1165	1244	5196	39.8%	42.2%
2014	10000	2678	2576	12678	12576	2906	3146	12678	40.6%	43.5%
2015	10000	342	95	10342	10095	698	1182	10342	13.0%	21.6%
2016	10000	3742	3949	13742	13949	4245	4245	13949	51.6%	51.6%
2017	1000	372	137	1372	1137	422	422	1372	52.1%	52.1%
2018	4700	456	201	5156	4901	1368	1368	5156	46.0%	46.0%
2019	0	110	201	110	201	199	199	201	100.0%	100.0%

1012 *Note: “Outplants” indicate the number of sockeye translocated into the Yakima River, and “Prosser” and “Roza” dam counts were*
 1013 *summed (“O+P” or “O+R”) with the outplants to provide an estimate of the total escapement of spawners in the Yakima River each*
 1014 *year. We used whichever number was greatest (“O+P” or “O+R”) to provide the maximum escapement of the Yakima River (“Max*
 1015 *YR”). A portion of these spawners were successfully genotyped either using 383 or 88 (the legacy panel) SNPs and tag rates were*
 1016 *calculated using the Max YR as the denominator. When cross information and the gender of the broodstock samples is unknown, then*
 1017 *the tag rate for single parentage is : $1-(f_t)^2$, where f_t is the fraction of the total broodstock not genotyped.*

1018 *Parentage based tagging assignments of Chinook salmon in harvest mixtures*

1019 A summary of the Chinook harvest samples that were genotyped (derived from Table 6)
1020 is presented in Table 24. Of the 3,993 harvest Chinook analyzed, PBT identified 1,944 hatchery-
1021 origin individuals that could be confidently assigned back to 79 hatchery broodstock sources
1022 (i.e., 3 Lower Columbia, 25 Snake River, and 51 Columbia River hatchery broodstocks) spawned
1023 in 2012-2016. The majority of PBT assigned individuals were from the 2014 brood year (i.e., 4-
1024 years-old).

1025 **Table 24. Summary of the Chinook salmon harvest samples by fishery, region, and fin clip**
1026 **in 2018.**

			Management period			Analysis			
Fishery	Region	Clip	Spring	Summer	Fall	GSI	PBT	Total	%PBT
Sport	A	AD	347			157	190	347	54.8%
	B	AD	526			273	253	526	48.1%
Test		AD	567			283	284	567	50.1%
	B	AI	83			58	25	83	30.1%
Ceremonial	Below_BON	AD	235			79	156	235	66.4%
		AI	38			25	13	38	34.2%
	ZN6	AD	139			50	89	139	64.0%
		AI	29			16	13	29	44.8%
Sport	A	AD		30		12	18	30	60.0%
	B	AD		42		13	29	42	69.0%
Treaty	ZN6	AD		165		31	134	165	81.2%
		AI		58		44	14	58	24.1%
Commercial	A	AD			198	53	145	198	73.2%
		AI			100	81	19	100	19.0%
Sport	A	AD			45	17	28	45	62.2%
		AI			119	94	25	119	21.0%
	B	AD			32	12	20	32	62.5%
		AI			79	67	12	79	15.2%
Test	B	AD			299	115	184	299	61.5%
		AI			172	147	25	172	14.5%
Treaty	ZN6	AD			234	44	190	234	81.2%
		AI			456	378	78	456	17.1%
						2049	1944	3993	48.7%

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Comparison of proportion of PBT assignments among Chinook salmon fisheries

The average of adipose-clipped Chinook salmon from fishery mixtures across all periods was that were assigned via PBT was 64% (range = 48% – 81%, Table 24). The fishery(s) with the minimum and maximum assigned adipose-clipped Chinook salmon was the spring sport fishery in region B and both summer and fall Treaty zone 6 fisheries, respectively. Among the adipose-intact fish, the average assignment via PBT was expectedly lower (average = 24%, range = 14% – 45%; Table 24). Among the fisheries with adipose-intact fish, the minimum and maximum PBT-assigned Chinook salmon was observed in the fall test and the spring ceremonial fishery, respectively. For both adipose-clipped and adipose-intact fish, the collections with the minimum PBT-assigned fish were due to areas that receive higher proportions of lower river hatchery fish that are not as well covered in the PBT baseline.

The test fishery in the Chinook Salmon Spring Management Period of 2018

We examined one source of information that could potentially be useful to managers particularly on years when the spring Chinook Salmon run is delayed. There is a test fishery that is typically conducted by WDFW on Sundays each week in the early spring (February – May). Four boats are contracted to perform a series of drifts (typically four drifts per boat) through the commercial zones 1-3 near the mouth of the Columbia River. The drifts are targeted for Spring Chinook and biodata and a tissue for genetic analysis is collected. Visual Stock ID classifies fish as lower river (West Cascade Spring and Willamette River stocks) versus upriver (all stocks destined above Bonneville Dam). Genetic Analysis can refine the initial Visual Stock ID calls and more accurately classify these fish into lower and upriver stocks. For example, we characterized reporting groups among the hatchery clipped, hatchery unclipped, and natural-origin VSI-upriver fish caught in the test fishery (Table 25, **Table 26**). We use a catch per unit effort (CPUE) metric in the test fishery that is based on the number of upriver Chinook handled per test fishery drift. If the test fishery CPUE were lagged 13 days the peak in CPUE appears to correspond with a peak in Bonneville weekly counts (Figure 21). So if the in-season genetic analysis includes the data from the test fishery, our report may be able to predict the abundance of particular hatchery and natural-origin stocks that we can expect to pass Bonneville Dam up to 2 weeks later which would be useful information for USvOR managers. In fact, comparison of the relative proportions of the hatchery broodstocks that were estimated in the test fishery and Bonneville Dam showed that these samples from these two sources have very similar compositions of stocks (Figure 19). Further, there is high correlation between the estimated CPUE of each hatchery broodstock in the test fishery and the estimated abundance of the same broodstocks passing Bonneville Dam in weeks lagged 13 days after the test fishery.

Table 25. Summary of the stock composition at the reporting group level of the VSI-identified upriver spring Chinook salmon in the test fishery of 2018 in units of catch-per-unit-effort (CPUE, # of fish per # of drifts).

Run type	Reporting Group Code	Hatchery origin-Clipped	Hatchery origin-No Clip	Natural origin- No Clip
		Estimated abundance	Estimated abundance	Estimated abundance

		Mea n	95% CI	Mean	95% CI	Mea n	95% CI
Spring	01_YOUNGS						
Spring	02_WCASSP	0.29	0.08 – 0.57				
Fall	03_WCASFA	0.04	0 – 0.15				
Spring	04_WILLAM	0.58	0.26 – 1				
Fall	05_SPCRTU						
Spring	06_KLICKR	0.04	0 – 0.15				
Spring	07_DESCSP	0.12	0 – 0.31			0.15	0.03 – 0.31
Spring	08_JOHNDR					0.04	0 – 0.09
Spring	09_YAKIMA	0.17	0 – 0.41			0.20	0.03 – 0.4
Spring	10_UCOLSP	0.61	0.22 – 1.07	0.13	0 – 0.4	0.23	0.06 – 0.44
Spring	11_TUCANO			0.08	0 – 0.24		
Spring/Summe r	12_HELLSC	2.37	1.74 – 2.98	0.28	0 – 0.58	0.24	0.07 – 0.45
Spring/Summe r	13_SFSALM	0.09	0 – 0.3				
Spring/Summe r	14_CHMBLN						
Spring/Summe r	15_MFSALM						
Spring/Summe r	16_UPSALM	0.26	0.07 – 0.53				
Fall	17_DESCFA						
Summer/Fall	18_UCOLSF						
Fall	19_SRFALL						
Spring	20_BONPOOLSP	2.05	1.38 – 2.66	0.13	0 – 0.39		
Spring	21_UMATILLASP	0.54	0.24 – 0.91	0.07	0 – 0.21		
Fall	22_BONPOOLFA						
Fall	23_UMATILLAFA						
	Total	7.15		0.70		0.86	

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Table 26. Summary of the stock composition at the broodstock level of the hatchery-origin VSI-identified upriver spring Chinook salmon in the test fishery of 2018 in units of catch-per-unit-effort (CPUE, # of fish per # of drifts).

Expected Run Time	Hatchery	code	Brood year	Adult Chinook AD			Adult Chinook AI			GSI RepGrp
				ML E	95% CI	Perc ent	ML E	95% CI	Perc ent	
01Spring	Parkdale Fish Facility	OtsPFF14_sp	2014	0.08	0 – 0.2	1.1%			0.0%	02_WCASSP
01Spring	Klickitat Hatchery	OtsKH14_sp	2014	0.04	0 – 0.12	0.5%			0.0%	06_KLICKR
01Spring	Round Butte Fish Hatchery	OtsRB14_sp	2014	0.08	0 – 0.19	1.1%			0.0%	07_DESCSP
01Spring	Yakima River Roza Dam	OtsYR14int_sp	2014	0.11	0 – 0.23	1.6%			0.0%	09_YAKIMA
01Spring	Yakima River Roza Dam	OtsYR14seg_sp	2014	0.06	0 – 0.17	0.8%			0.0%	09_YAKIMA
01Spring	Chief Joseph Hatchery	OtsCJH14_sp	2014	0.04	0 – 0.13	0.6%			0.0%	10_UCOLSP
01Spring	Leavenworth National Fish Hatchery	OtsLNFH14_sp	2014	0.08	0 – 0.15	1.1%			0.0%	10_UCOLSP
01Spring	Winthrop National Fish Hatchery	OtsWTP14_sp	2014	0.15	0.04 – 0.28	2.1%	0.13	0 – 0.31	18.5%	10_UCOLSP
01Spring	Winthrop National Fish Hatchery	OtsWTP15_sp	2015	0.04	0 – 0.11	0.5%			0.0%	10_UCOLSP
01Spring	Lyons Ferry Fish Hatchery	OtsTUCW14S_sp	2014			0.0%	0.08	0 – 0.16	11.5%	11_TUCANO
01Spring	Clearwater Fish Hatchery	OtsCLWH14S_sp	2014	0.66	0.43 – 0.89	9.3%			0.0%	12_HELLSC
01Spring	Clearwater Fish Hatchery	OtsCLWH15S_sp	2015	0.04	0 – 0.11	0.5%			0.0%	12_HELLSC
01Spring	Clearwater Fish Hatchery	OtsPOWP14S_sp	2014	0.38	0.19 – 0.57	5.3%	0.12	0 – 0.29	17.5%	12_HELLSC
01Spring	Dworshak National Fish Hatchery	OtsDWOR15S_sp	2015	0.08	0 – 0.2	1.1%	0.04	0 – 0.12	5.6%	12_HELLSC

01Spring	Lookingglass Fish Hatchery	OtsCTHW14S_sp	2014	0.0 4	0 – 0.13	0.6%			0.0%	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsGRUW14S_sp	2014	0.0 4	0 – 0.12	0.5%	0.09	0 – 0.26	12.3 %	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsLOOK14S_sp	2014	0.0 8	0 – 0.17	1.1%			0.0%	12_HELLSC
01Spring	Nez Perce	OtsNPFH14S_sp	2014			0.0%	0.04	0 – 0.11	5.3%	12_HELLSC
01Spring	Carson National Fish Hatchery	OtsCAR13_sp	2013	0.0 5	0 – 0.14	0.7%			0.0%	20_BONPOO LSP
01Spring	Carson National Fish Hatchery	OtsCAR14_sp	2014	0.6 1	0.38 – 0.86	8.5%	0.05	0 – 0.1	6.9%	20_BONPOO LSP
01Spring	Little White Salmon National Fish Hatchery	OtsLWS13_sp	2013	0.0 6	0 – 0.14	0.9%			0.0%	20_BONPOO LSP
01Spring	Little White Salmon National Fish Hatchery	OtsLWS14_sp	2014	1.3 3	1 – 1.65	18.6 %	0.08	0 – 0.25	12.1 %	20_BONPOO LSP
01Spring	Umatilla Fish Hatchery	OtsUMA14_s p	2014	0.5 0	0.3 – 0.74	7.0%	0.07	0 – 0.14	10.3 %	21_UMATIL LASP
01Spring	Umatilla Fish Hatchery	OtsUMA15_s p	2015	0.0 4	0 – 0.12	0.5%			0.0%	21_UMATIL LASP
02Spring/Summer	McCall Fish Hatchery	OtsMCCA14S_spsu	2014	0.0 5	0 – 0.16	0.7%			0.0%	13_SFSALM
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT14S_spsu	2014	0.2 6	0.11 – 0.45	3.7%			0.0%	16_UPSALM
#N/A	#N/A	Unassigned	#N/A	2.2 6	1.85 – 2.66	31.6 %			0.0%	#N/A
		TOTAL		7.1 5		100.0 %	0.70		100.0 %	

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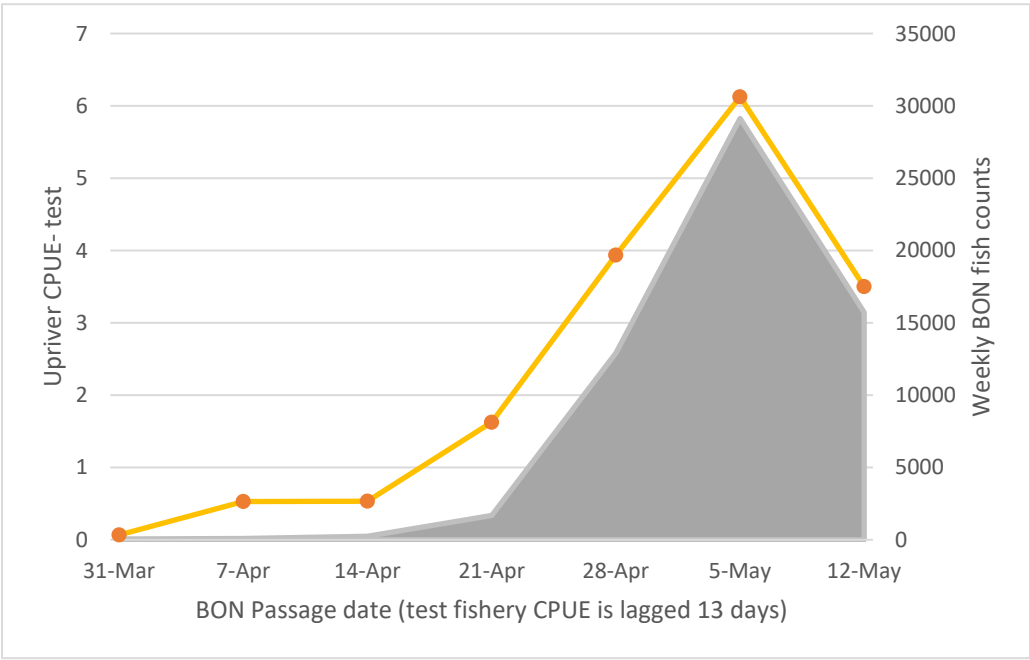


Figure 18. The relationship between the test fishery upriver Chinook Salmon CPUE (line) and weekly fish counts at Bonneville Dam (solid gray) in 2018.

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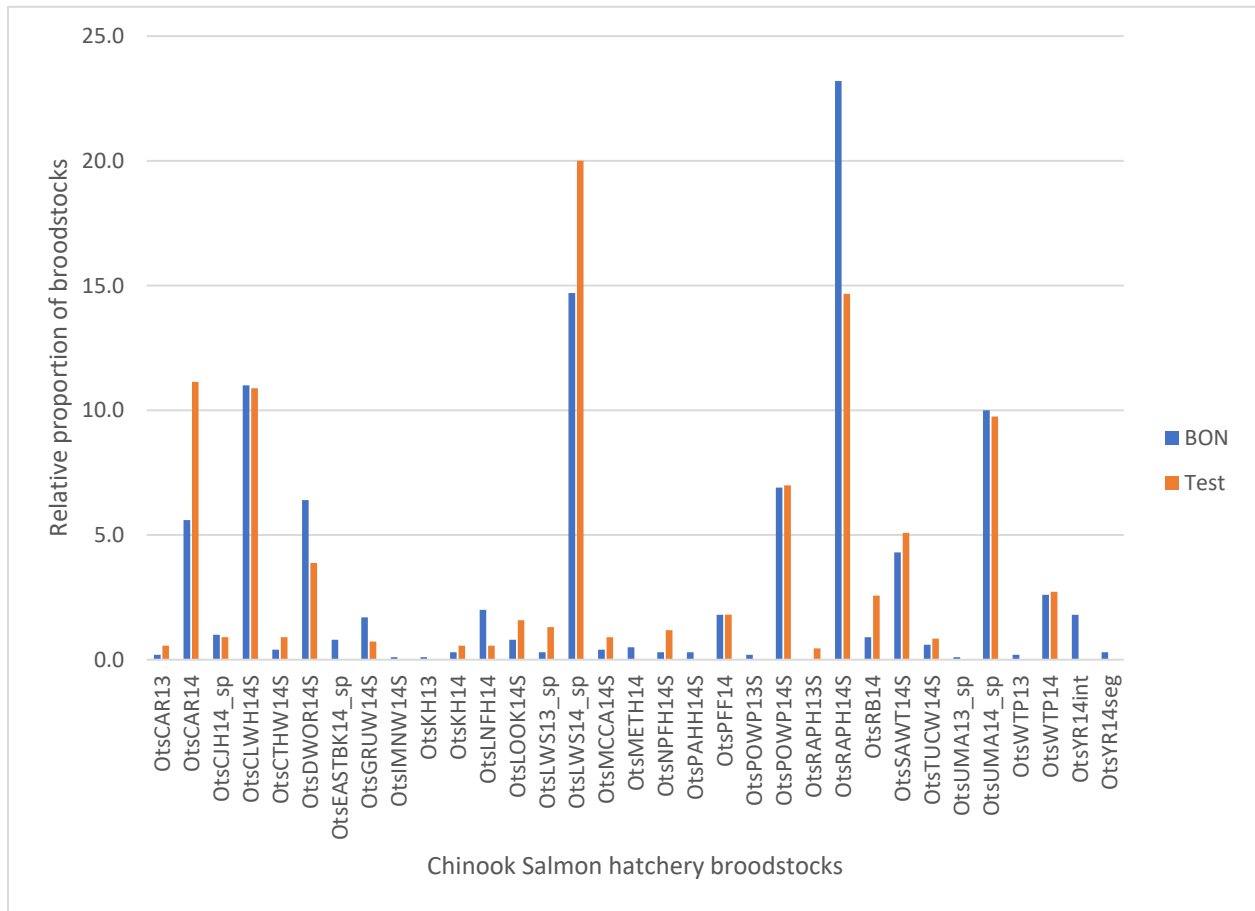


Figure 19. Relative proportions of hatchery broodstocks of upriver Chinook Salmon caught in the test fishery (March 18 – April 29) compared to Chinook Salmon that passed Bonneville Dam in corresponding weeks lagged 13 days from the test fishery (March 31 – May 12) in 2018.

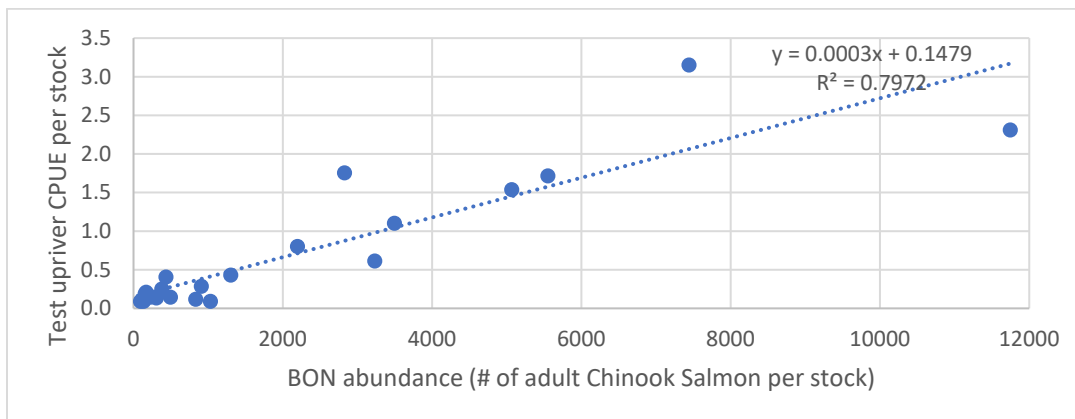


Figure 20. Correlation of the test fishery CPUE with the estimated abundance passing Bonneville Dam of hatchery broodstocks identified by PBT analysis in 2018.

We reported spring sport fishery results for reporting group level stock composition (Table 27) and the hatchery broodstock composition (Table 28) of the kept fish (clipped hatchery-origin fish).

Table 27. Summary of the stock composition at the reporting group level of the kept spring Chinook salmon in the Non-Treaty sport fishery of 2018 in units of reported catch.

Run type	Reporting Group Code	Hatchery origin-Clipped	
		Estimated abundance	
		Mean	95% CI
Spring	01_YOUNGS		
Spring	02_WCASSP	675	489 – 871
Fall	03_WCASFA	36	0 – 91
Spring	04_WILLAM	1,133	905 – 1368
Fall	05_SPCRTU		
Spring	06_KLICKR	94	29 – 170
Spring	07_DESCSP	60	9 – 122
Spring	08_JOHNDR		
Spring	09_YAKIMA	96	24 – 199
Spring	10_UCOLSP	459	296 – 634
Spring	11_TUCANO		
Spring/Summer	12_HELLSC	1,884	1607 – 2186
Spring/Summer	13_SFSALM	369	214 – 549
Spring/Summer	14_CHMBLN		
Spring/Summer	15_MFSALM		
Spring/Summer	16_UPSALM	80	19 – 159
Fall	17_DESCFA		
Summer/Fall	18_UCOLSF	718	558 – 858
Fall	19_SRFALL	7	0 – 44
Spring	20_BONPOOLSP	1,579	1299 – 1829
Spring	21_UMATILLASP	319	201 – 447
Fall	22_BONPOOLFA		
Fall	23_UMATILLAFA		
Total		7,509	

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Table 28. Summary of the stock composition at the broodstock level of the kept spring Chinook salmon in the Non-Treaty sport fishery of 2018 in units of reported catch.

Adult Chinook AD							
Expected Run Time	Hatchery	Broodstock	Broodyear	MLE	95% CI	Percent	GSI RepGrp
01Spring	Parkdale Fish Facility	OtsPFF13_sp	2013	13	0 – 39	0.2%	02_WCASSP
01Spring	Parkdale Fish Facility	OtsPFF14_sp	2014	104	59 – 161	1.4%	02_WCASSP
01Spring	Klickitat Hatchery	OtsKH13_sp	2013	37	9 – 71	0.5%	06_KLICKR
01Spring	Klickitat Hatchery	OtsKH14_sp	2014	29	9 – 59	0.4%	06_KLICKR
01Spring	Round Butte Fish Hatchery	OtsRB14_sp	2014	47	18 – 84	0.6%	07_DESCSP
01Spring	Round Butte Fish Hatchery	OtsRB15_sp	2015	12	0 – 37	0.2%	07_DESCSP
01Spring	Yakima River Roza Dam	OtsYR14seg_sp	2014	71	27 – 125	0.9%	09_YAKIMA
01Spring	Yakima River Roza Dam	OtsYR14int_sp	2014	20	0 – 43	0.3%	09_YAKIMA
01Spring	Chief Joseph Hatchery	OtsCJH14_sp	2014	41	10 – 82	0.5%	10_UCOLSP
01Spring	Eastbank Fish Hatchery	OtsEASTBK14_sp	2014	18	0 – 37	0.2%	10_UCOLSP
01Spring	Eastbank Fish Hatchery	OtsEASTBK15_sp	2015	23	0 – 46	0.3%	10_UCOLSP
01Spring	Leavenworth National Fish Hatchery	OtsLNFH14_sp	2014	64	28 – 110	0.9%	10_UCOLSP
01Spring	Winthrop National Fish Hatchery	OtsWTP14_sp	2014	29	0 – 59	0.4%	10_UCOLSP
01Spring	Clearwater Fish Hatchery	OtsCLWH14S_sp	2014	476	368 – 587	6.3%	12_HELLSC
01Spring	Clearwater Fish Hatchery	OtsPOWP14S_sp	2014	168	110 – 232	2.2%	12_HELLSC
01Spring	Dworshak National Fish Hatchery	OtsDWOR15S_sp	2015	12	0 – 35	0.2%	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsCTHW14S_sp	2014	32	10 – 63	0.4%	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsLOOK14S_sp	2014	62	29 – 104	0.8%	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsLSTW14S_sp	2014	30	0 – 74	0.4%	12_HELLSC
01Spring	Carson National Fish Hatchery	OtsCAR13_sp	2013	44	11 – 89	0.6%	20_BONPOOLSP
01Spring	Carson National Fish Hatchery	OtsCAR14_sp	2014	625	504 – 751	8.3%	20_BONPOOLSP
01Spring	Little White Salmon National Fish Hatchery	OtsLWS13_sp	2013	54	18 – 91	0.7%	20_BONPOOLSP
01Spring	Little White Salmon National Fish Hatchery	OtsLWS14_sp	2014	856	718 – 994	11.4%	20_BONPOOLSP
01Spring	Umatilla Fish Hatchery	OtsUMA13_sp	2013	9	0 – 28	0.1%	21_UMATILLASP
01Spring	Umatilla Fish Hatchery	OtsUMA14_sp	2014	309	227 – 398	4.1%	21_UMATILLASP

02Spring/Summer	McCall Fish Hatchery	OtsMCCA14S_spsu	2014	307	212 – 416	4.1%	13_SFSALM
02Spring/Summer	McCall Fish Hatchery	OtsMCCA15S_spsu	2015	25	0 – 51	0.3%	13_SFSALM
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT14S_spsu	2014	31	9 – 61	0.4%	16_UPSALM
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT15S_spsu	2015	11	0 – 33	0.1%	16_UPSALM
03Summer	Chief Joseph Hatchery	OtsCJH13int_su	2013	16	0 – 49	0.2%	18_UCOLSF
03Summer	Chief Joseph Hatchery	OtsCJH14int_su	2014	76	38 – 127	1.0%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK12_su	2012	34	11 – 68	0.5%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK13_su	2013	45	11 – 80	0.6%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK14_su	2014	90	45 – 136	1.2%	18_UCOLSF
03Summer	Entiat National Fish Hatchery	OtsENFH14_su	2014	92	31 – 154	1.2%	18_UCOLSF
03Summer	Wells Fish Hatchery	OtsWELLS13_su	2013	55	21 – 98	0.7%	18_UCOLSF
03Summer	Wells Fish Hatchery	OtsWELLS14_su	2014	113	57 – 170	1.5%	18_UCOLSF
#N/A	#N/A	Unassigned	#N/A	3428	3191 – 3669	45.7%	
		TOTAL		7,509		100.0%	

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Our analyses of the spring Treaty permit fishery below Bonneville Dam, allowed us to estimate the portion of stocks that were natural-origin (8.8%) compared to the total catch that included clipped and unclipped hatchery-origin Chinook salmon (Table 29). We could further estimate abundances of hatchery broodstocks among the hatchery-origin fish (Table 30).

Table 29. Summary of the stock composition at the reporting group level of the Yakama Nation permit fishery below Bonneville Dam in 2018 in units of reported catch.

Reporting Group name	Run type	Reporting Group Code	Hatchery origin- Clipped		Hatchery origin- No Clip		Natural origin- No Clip	
			Estimated abundance		Estimated abundance		Estimated abundance	
			Mean	95% CI	Mean	95% CI	Mean	95% CI
Youngs Bay	Spring	01_YOUNGS						
West Cascade Spring	Spring	02_WCASSP	7	0 – 16				
West Cascade Fall	Fall	03_WCASFA						
Willamette	Spring	04_WILLAM						
Spring Creek Tule	Fall	05_SPCRTU						
Klickitat	Spring	06_KLICKR	2	0 – 7				
Deschutes spring	Spring	07_DESCSP	8	2 – 18			2	0 – 5
John Day	Spring	08_JOHNDR	2	0 – 7			3	0 – 7
Yakima	Spring	09_YAKIMA	5	0 – 15			2	0 – 5
Upper Columbia spring	Spring	10_UCOLSP	41	22 – 64	5	0 – 14	9	3 – 16
Tucannon	Spring	11_TUCANO			4	0 – 11		
Hells Canyon	Spring/Summer	12_HELLSC	192	156 – 225	10	2 – 24	13	6 – 21
South Fork Salmon	Spring/Summer	13_SFSALM	5	0 – 13			2	0 – 5
Chamberlain Creek	Spring/Summer	14_CHMBLN						
Middle Fork Salmon	Spring/Summer	15_MFSALM					3	0 – 7
Upper Salmon	Spring/Summer	16_UPSALM	18	7 – 34			7	2 – 13

Deschutes fall	Fall	17_DESCFA						
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF						
Snake River fall	Fall	19_SRFALL						
Bonneville Pool spring	Spring	20_BONPOOLSP	84	57 – 111	4	0 – 12		
Umatilla spring	Spring	21_UMATILLASP	28	12 – 46				
Bonneville Pool fall	Fall	22_BONPOOLFA						
Umatilla fall	Fall	23_UMATILLAFA						
Total			391		23		40	

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1113 **Table 30. Summary of the stock composition at the broodstock level of the Yakama Nation permit fishery below**
1114 **Bonneville Dam of 2018 in units of reported catch.**

Expected Run Time	Hatchery	Broodstock	Broody ear	Adult Chinook AD			Adult Chinook AI			GSI RepGrp
				ML E	95% CI	Perce nt	ML E	95% CI	Perce nt	
01Spring	Parkdale Fish Facility	OtsPFF14_sp	2014	7	2 – 14	1.8%			0.0%	02_WCASSP
01Spring	Round Butte Fish Hatchery	OtsRB14_sp	2014	8	3 – 15	2.1%			0.0%	07_DESCSP
01Spring	Yakima River Roza Dam	OtsYR14int_s p	2014	5	2 – 10	1.3%			0.0%	09_YAKIMA
01Spring	Chief Joseph Hatchery	OtsCJH15_sp	2015			0.0%	2	0 – 5	7.3%	10_UCOLSP
01Spring	Eastbank Fish Hatchery	OtsEASTBK1 4_sp	2014	5	0 – 10	1.3%			0.0%	10_UCOLSP
01Spring	Leavenworth National Fish Hatchery	OtsLNFH14_s p	2014	8	3 – 15	2.1%			0.0%	10_UCOLSP
01Spring	Winthrop National Fish Hatchery	OtsWTP14_sp	2014	7	2 – 12	1.8%	4	0 – 9	15.6 %	10_UCOLSP
01Spring	Lyons Ferry Fish Hatchery	OtsTUCW14S _sp	2014			0.0%	4	0 – 9	16.3 %	11_TUCANO
01Spring	Clearwater Fish Hatchery	OtsCLWH14S _sp	2014	60	43 – 76	15.4 %			0.0%	12_HELLSC
01Spring	Clearwater Fish Hatchery	OtsPOWP14S _sp	2014	22	12 – 32	5.6%	3	0 – 7	14.8 %	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsCTHW14S _sp	2014	4	0 – 8	1.0%	2	0 – 6	8.4%	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsGRUW14S _sp	2014	2	0 – 5	0.4%	3	0 – 8	14.7 %	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsLOOK14S _sp	2014	9	4 – 16	2.3%			0.0%	12_HELLSC
01Spring	Lookingglass Fish Hatchery	OtsCTHW15S _sp	2015	2	0 – 5	0.4%			0.0%	12_HELLSC
01Spring	Nez Perce	OtsNPFH14S_ sp	2014			0.0%	2	0 – 5	7.5%	12_HELLSC
01Spring	Rapid River Fish Hatchery	OtsRAPH15S _sp	2015	2	0 – 5	0.5%			0.0%	12_HELLSC

01Spring	Carson National Fish Hatchery	OtsCAR14_sp	2014	35	24 – 49	9.1%	2	0 – 6	8.2%	20_BONPOO LSP
01Spring	Carson National Fish Hatchery	OtsCAR15_sp	2015	2	0 – 5	0.4%			0.0%	20_BONPOO LSP
01Spring	Little White Salmon National Fish Hatchery	OtsLWS14_sp	2014	46	31 – 60	11.9 %	2	0 – 5	7.3%	20_BONPOO LSP
01Spring	Umatilla Fish Hatchery	OtsUMA14_s p	2014	28	18 – 40	7.2%			0.0%	21_UMATIL LASP
02Spring/Summer	McCall Fish Hatchery	OtsMCCA14S_spsu	2014	2	0 – 7	0.6%			0.0%	13_SFSALM
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT14S_spsu	2014	13	7 – 22	3.4%			0.0%	16_UPSALM
#N/A	#N/A	Unassigned	#N/A	123	101 – 146	31.5 %			0.0%	#N/A
		TOTAL		391		100.0 %	23		100.0 %	

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For the summer Non-Treaty sport fishery below Bonneville Dam, we estimated report group composition (Table 31) of the kept fish (clipped hatchery-origin), and further broke these groups into abundance estimates of broodstocks (Table 32).

Table 31. Summary of the stock composition at the reporting group level of the summer Non-Treaty sport fishery below Bonneville Dam in 2018 in units of reported catch.

Reporting Group name	Run type	Reporting Group Code	Hatchery origin- Clipped	
			Estimated abundance	
			Mean	95% CI
Youngs Bay	Spring	01_YOUNGS		
West Cascade Spring	Spring	02_WCASSP	14	0 – 56
West Cascade Fall	Fall	03_WCASFA		
Willamette	Spring	04_WILLAM	180	56 – 309
Spring Creek Tule	Fall	05_SPCRTU		
Klickitat	Spring	06_KLICKR		
Deschutes spring	Spring	07_DESCSP		
John Day	Spring	08_JOHNDR		
Yakima	Spring	09_YAKIMA		
Upper Columbia spring	Spring	10_UCOLSP		
Tucannon	Spring	11_TUCANO		
Hells Canyon	Spring/Summer	12_HELLSC	17	0 – 66
South Fork Salmon	Spring/Summer	13_SFSALM	40	0 – 119
Chamberlain Creek	Spring/Summer	14_CHMBLN		
Middle Fork Salmon	Spring/Summer	15_MFSALM		
Upper Salmon	Spring/Summer	16_UPSALM	22	0 – 83
Deschutes fall	Fall	17_DESCFA		
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	755	627 – 925
Snake River fall	Fall	19_SRFALL		
Bonneville Pool spring	Spring	20_BONPOOLSP		
Umatilla spring	Spring	21_UMATILLASP		
Bonneville Pool fall	Fall	22_BONPOOLFA		
Umatilla fall	Fall	23_UMATILLAFA		
Total			1,027	

1127 **Table 32. Summary of the stock composition at the broodstock level of the summer Non-Treaty sport fishery below**
 1128 **Bonneville Dam of 2018 in units of reported catch.**

Summer Sport Fishery 2018				Adult Chinook AD			
Expected Run Time	Hatchery	Broodstock	Broodyear	MLE	95% CI	Percent	GSI RepGrp
01Spring	South Santiam Hatchery	OtsSSANT16_sp	2016	15	0 – 45	1.5%	04_WILLAM
02Spring/Summer	Lookingglass Fish Hatchery	OtsIMNW15S_spsu	2015	17	0 – 50	1.6%	12_HELLSC
02Spring/Summer	McCall Fish Hatchery	OtsMCCA14S_spsu	2014	40	0 – 79	3.9%	13_SFSALM
03Summer	Chief Joseph Hatchery	OtsCJH13seg_su	2013	48	16 – 96	4.7%	18_UCOLSF
03Summer	Chief Joseph Hatchery	OtsCJH13int_su	2013	61	20 – 123	6.0%	18_UCOLSF
03Summer	Chief Joseph Hatchery	OtsCJH14int_su	2014	128	64 – 208	12.5%	18_UCOLSF
03Summer	Chief Joseph Hatchery	OtsCJH14seg_su	2014	65	0 – 162	6.3%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK12_su	2012	29	0 – 57	2.8%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK13_su	2013	57	14 – 100	5.6%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK14_su	2014	143	71 – 214	13.9%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK15_su	2015	14	0 – 43	1.4%	18_UCOLSF
03Summer	Entiat National Fish Hatchery	OtsENFH14_su	2014	136	58 – 214	13.3%	18_UCOLSF
03Summer	Wells Fish Hatchery	OtsWELLS13_su	2013	14	0 – 43	1.4%	18_UCOLSF
03Summer	Wells Fish Hatchery	OtsWELLS14_su	2014	29	0 – 71	2.8%	18_UCOLSF
#N/A	#N/A	Unassigned	#N/A	232	120 – 346	22.6%	#N/A
		TOTAL		1,027		100.0%	

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1130 The stock composition of the summer Treaty zone 6 commercial fishery was characterized by reporting group level of the
 1131 clipped and unclipped hatchery-origin and the natural-origin Chinook salmon (Table 33). We further categorized hatchery-origin
 1132 stocks by broodstock (**Table 34**).
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1134 **Table 33. Summary of the stock composition at the reporting group level of the summer Treaty Zone 6 commercial fishery in**
 1135 **2018 in units of reported catch.**

Run type	Reporting Group Code	Hatchery origin- Clipped		Hatchery origin- No Clip		Natural origin- No Clip	
		Estimated abundance		Estimated abundance		Estimated abundance	
		Mean	95% CI	Mean	95% CI	Mean	95% CI
Spring	01_YOUNGS						
Spring	02_WCASSP						
Fall	03_WCASFA						
Spring	04_WILLAM	35	0 – 170			28	0 – 85
Fall	05_SPCRTU						
Spring	06_KLICKR						
Spring	07_DESCSP					31	0 – 91
Spring	08_JOHNDR						
Spring	09_YAKIMA						
Spring	10_UCOLSP	36	0 – 249	48	0 – 193	28	0 – 85
Spring	11_TUCANO						
Spring/Summer	12_HELLSC	397	58 – 815	33	0 – 131	28	0 – 85
Spring/Summer	13_SFSALM	291	0 – 730	179	0 – 403	80	6 – 179
Spring/Summer	14_CHMBLN						
Spring/Summer	15_MFSALM						
Spring/Summer	16_UPSALM	105	0 – 419			49	0 – 125
Fall	17_DESCFA						
Summer/Fall	18_UCOLSF	7,226	6573 – 7710	271	68 – 545	961	748 – 1158
Fall	19_SRFALL					28	0 – 85
Spring	20_BONPOOLSP						

Spring	21_UMATILLASP						
Fall	22_BONPOOLFA						
Fall	23_UMATILLAFA						
	Total	8,090		531		1,234	

Table 34. Summary of the stock composition at the broodstock level of the summer Treaty Zone 6 commercial fishery of 2018 in units of reported catch.

Summer Treaty Fishery 2018					Adult Chinook AD		Adult Chinook AI			
Expected Run Time	Hatchery	Broodstock	Broody ear	ML E	95% CI	Percent	ML E	95% CI	Percent	GSI RepGrp
01Spring	Eastbank Fish Hatchery	OtsEASTBK14_sp	2014			0.0%	48	0 – 145	9.1%	10_UCOL SP
01Spring	Clearwater Fish Hatchery	OtsCLWH14S_sp	2014	54	0 – 163	0.7%			0.0%	12_HELL SC
01Spring	Clearwater Fish Hatchery	OtsPOWP14S_sp	2014			0.0%	33	0 – 98	6.2%	12_HELL SC
01Spring	Lookingglass Fish Hatchery	OtsLSTW14S_sp	2014	342	124 – 595	4.2%			0.0%	12_HELL SC
02Spring/Summer	McCall Fish Hatchery	OtsMCCA14S_spsu	2014	291	73 – 528	3.6%	179	45 – 313	33.7 %	13_SFSA LM
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT14S_spsu	2014	105	0 – 262	1.3%			0.0%	16_UPSA LM
03Summer	Chief Joseph Hatchery	OtsCJH13seg_su	2013	645	364 – 920	8.0%	32	0 – 96	6.0%	18_UCOL SF
03Summer	Chief Joseph Hatchery	OtsCJH13int_su	2013	185	0 – 350	2.3%			0.0%	18_UCOL SF
03Summer	Chief Joseph Hatchery	OtsCJH14int_su	2014	914	592 – 1265	11.3 %	36	0 – 108	6.8%	18_UCOL SF
03Summer	Chief Joseph Hatchery	OtsCJH14seg_su	2014	766	314 – 1180	9.5%			0.0%	18_UCOL SF
03Summer	Eastbank Fish Hatchery	OtsEASTBK12su	2012	154	45 – 309	1.9%			0.0%	18_UCOL SF

03Summer	Eastbank Fish Hatchery	OtsEASTBK13_su	2013	332	121 – 553	4.1%	75	0 – 149	14.1%	18_UCOLSF
03Summer	Eastbank Fish Hatchery	OtsEASTBK14_su	2014	1,340	930 – 1722	16.6%	99	30 – 199	18.7%	18_UCOLSF
03Summer	Entiat National Fish Hatchery	OtsENFH13_su	2013	58	0 – 174	0.7%			0.0%	18_UCOLSF
03Summer	Entiat National Fish Hatchery	OtsENFH14_su	2014	1,422	959 – 1850	17.6%			0.0%	18_UCOLSF
03Summer	Entiat National Fish Hatchery	OtsENFH15_su	2015	59	0 – 177	0.7%			0.0%	18_UCOLSF
03Summer	Wells Fish Hatchery	OtsWELLS13_su	2013	401	186 – 641	5.0%			0.0%	18_UCOLSF
03Summer	Wells Fish Hatchery	OtsWELLS14_su	2014	929	604 – 1276	11.5%	29	0 – 87	5.5%	18_UCOLSF
#N/A	#N/A	Unassigned	#N/A	92	0 – 682	1.1%			0.0%	#N/A
		TOTAL		8,090		100.0%	531		100.0%	

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1142 The fall test (pound net) fishery reported a total of 600 kept Chinook salmon (all clipped hatchery-origin), which were
 1143 comprised of a third lower river stocks (primarily 03_WCASFA), and 2/3 upriver stocks (18% Spring Creek tules and the remaining
 1144 were upriver brights). Among the released stocks, 25% were lower river and the rest were upriver stocks (Table 35). The hatchery-
 1145 origin fish were identified by broodstock (**Table 36**), and the lower river stocks were found to be comprised of Big Creek, Cowlitz
 1146 Salmon Hatchery, North Toutle Hatchery, and Washougal Fish Hatchery.

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 1148 **Table 35. Summary of the stock composition at the reporting group level of the fall test fishery in 2018 in units of reported**
 1149 **catch.**

Reporting Group name	Run type	Reporting Group Code	RELEASED Chinook salmon					
			Hatchery-origin Kept- Clipped		Hatchery origin- No Clip		Natural origin- No Clip	
			Estimated abundance		Estimated abundance		Estimated abundance	
			Mean	95% CI	Mean	95% CI	Mean	95% CI
Youngs Bay	Spring	01_YOUNGS					21	5 – 38
West Cascade Spring	Spring	02_WCASSP						
West Cascade Fall	Fall	03_WCASFA	196	138 – 244			212	166 – 263
Willamette	Spring	04_WILLAM	2	0 – 7				
Spring Creek Tule	Fall	05_SPCRTU	109	72 – 166	28	3 – 59	45	23 – 71
Klickitat	Spring	06_KLICKR						
Deschutes spring	Spring	07_DESCSP						
John Day	Spring	08_JOHNDR						
Yakima	Spring	09_YAKIMA						
Upper Columbia spring	Spring	10_UCOLSP						
Tucannon	Spring	11_TUCANO						
Hells Canyon	Spring/Summer	12_HELLSC						
South Fork Salmon	Spring/Summer	13_SFSALM						
Chamberlain Creek	Spring/Summer	14_CHMBLN						
Middle Fork Salmon	Spring/Summer	15_MFSALM						
Upper Salmon	Spring/Summer	16_UPSALM						
Deschutes fall	Fall	17_DESCFA					5	0 – 15

Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	76	47 – 111	42	10 – 88	431	376 – 488
SNAKE RIVER fall	Fall	19_SRFALL	71	43 – 100	42	12 – 82	68	38 – 98
Bonneville Pool spring	Spring	20_BONPOOLSP						
UMATILLA spring	Spring	21_UMATILLASP						
Bonneville Pool fall	Fall	22_BONPOOLFA	139	101 – 176	9	0 – 28		
UMATILLA fall	Fall	23_UMATILLAFA	8	0 – 23				
Total			600		121		782	

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Table 36. Summary of the stock composition at the broodstock level of the fall test fishery of 2018 in units of reported catch.

Fall Pound Net (Test Fishery 2018)				Kept Chinook AD			Released Chinook AI			GSI RepGrp
Expected Run Time	Hatchery	Broodstock	Broody ear	ML E	95% CI	Percent	ML E	95% CI	Percent	
04Fall	Big Creek Hatchery	OtsBIG15_fa	2015	41	14 – 81	6.8%			0.0%	03_WCASFA
04Fall	Cowlitz Salmon Hatchery	OtsCOW15_fa	2015	2	0 – 7	0.4%			0.0%	03_WCASFA
04Fall	North Toutle Hatchery	OtsTOU15_fa	2015	10	2 – 17	1.6%			0.0%	03_WCASFA
04Fall	Washougal Fish Hatchery	OtsWAS15_fa	2015	10	0 – 30	1.7%			0.0%	03_WCASFA
04Fall	Spring Creek National Fish Hatchery	OtsSPCR15_fa	2015	85	65 – 107	14.2%	28	10 – 47	22.9%	05_SPCRTU
04Fall	Spring Creek National Fish Hatchery	OtsSPCR16_fa	2016	7	0 – 14	1.1%			0.0%	05_SPCRTU
04Fall	Priest Rapids Hatchery	OtsPRH13_fa	2013	2	0 – 5	0.3%	7	0 – 20	5.6%	18_UCOLSF
04Fall	Priest Rapids Hatchery	OtsPRH14_fa	2014	28	16 – 41	4.7%	9	0 – 20	7.2%	18_UCOLSF
04Fall	Priest Rapids Hatchery	OtsPRH15_fa	2015	29	17 – 42	4.8%	27	10 – 47	22.3%	18_UCOLSF
04Fall	Lyons Ferry Fish Hatchery	OtsLYON14S_1_fa	2014	27	15 – 39	4.5%	8	0 – 20	7.0%	19_SRFALL
04Fall	Lyons Ferry Fish Hatchery	OtsLYON15S_1_fa	2015	25	13 – 38	4.1%	16	5 – 33	13.5%	19_SRFALL
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH14S_1_fa	2014	8	2 – 16	1.4%	9	0 – 21	7.2%	19_SRFALL
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH15S_1_fa	2015	2	0 – 7	0.4%	9	0 – 21	7.2%	19_SRFALL
04Fall	Little White Salmon National Fish Hatchery	OtsLWS13_fa	2013	18	9 – 29	3.0%			0.0%	22_BONPOO LFA
04Fall	Little White Salmon National Fish Hatchery	OtsLWS14_fa	2014	45	31 – 61	7.5%	3	0 – 10	2.8%	22_BONPOO LFA
04Fall	Little White Salmon National Fish Hatchery	OtsLWS15_fa	2015	76	56 – 96	12.7%	5	0 – 16	4.4%	22_BONPOO LFA
04Fall	Umatilla Fish Hatchery	OtsUMA13_fa	2013	4	0 – 11	0.7%			0.0%	23_UMATILL AFA

04Fall	Umatilla Fish Hatchery	OtsUMA14_fa	2014	4	0 – 9	0.7%			0.0%	23_UMATILL AFA
#N/A	#N/A	Unassigned	#N/A	177	129 – 221	29.6%			0.0%	#N/A
		TOTAL		600		100.0 %	121		100.0 %	

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1155 The fall Non-Treaty sport fishery was not mark-selective and could be characterized by reporting group composition of clipped
 1156 and unclipped hatchery-origin and natural-origin stocks (**Table 37**). We also reported the broodstock composition of the hatchery-
 1157 origin Chinook salmon (**Table 38**).
 1158

1159 **Table 37. Summary of the stock composition at the reporting group level of the kept adult Chinook salmon of the fall**
 1160 **Non-Treaty sport fishery in 2018 in units of reported catch.**

Reporting Group name	Run type	Reporting Group Code	Hatchery origin- Clipped		Hatchery origin- No Clip		Natural origin- No Clip	
			Estimated abundance		Estimated abundance		Estimated abundance	
			Mea n	95% CI	Mean	95% CI	Mea n	95% CI
Youngs Bay	Spring	01_YOUNGS					84	0 – 170
West Cascade Spring	Spring	02_WCASSP						
West Cascade Fall	Fall	03_WCASFA	344	74 – 629	218	0 – 846	597	269 – 940
Willamette	Spring	04_WILLAM					42	0 – 126
Spring Creek Tule	Fall	05_SPCRTU	371	124 – 699				
Klickitat	Spring	06_KLICKR						
Deschutes spring	Spring	07_DESCSP						
John Day	Spring	08_JOHNDR						
Yakima	Spring	09_YAKIMA						
Upper Columbia spring	Spring	10_UCOLSP						
Tucannon	Spring	11_TUCANO						
Hells Canyon	Spring/Summe r	12_HELLSC						
South Fork Salmon	Spring/Summe r	13_SFSALM						
Chamberlain Creek	Spring/Summe r	14_CHMBLN						
Middle Fork Salmon	Spring/Summe r	15_MFSALM						
Upper Salmon	Spring/Summe r	16_UPSALM						

Deschutes fall	Fall	17_DESCFA					233	79 – 386
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	835	417 – 1321	413	121 – 753	3,674	3203 – 4172
Snake River fall	Fall	19_SRFALL	398	119 – 729	332	84 – 659	1,346	997 – 1705
Bonneville Pool spring	Spring	20_BONPOOLSP						
Umatilla spring	Spring	21_UMATILLASP						
Bonneville Pool fall	Fall	22_BONPOOLFA	482	159 – 842	397	116 – 760		
Umatilla fall	Fall	23_UMATILLAFA	37	0 – 186				
Total			2,467		1,360		5,975	

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Table 38. Summary of the stock composition at the broodstock level of the kept adult Chinook salmon of the fall Non-Treaty sport fishery in 2018 in units of reported catch.

Fall Sport Fishery 2018					Adult Chinook AD			Adult Chinook AI			
Ord er	Expected Run Time	Hatchery	Broodstock	Brood year	M LE	95% CI	Perc ent	M LE	95% CI	Perc ent	GSI RepGrp
	03Summer	Chief Joseph Hatchery	OtsCJH14se g_su	2014	95	0 – 286	3.9%			0.0%	18_UCOLSF
	04Fall	North Toutle Hatchery	OtsTOU15_f a	2015			0.0%	48	0 – 145	3.6%	03_WCASFA
	04Fall	Washougal Fish Hatchery	OtsWAS15_f a	2015			0.0%	169	0 – 508	12.4 %	03_WCASFA
	04Fall	Spring Creek National Fish Hatchery	OtsSPCR15_ fa	2015	298	130 – 476	12.1 %			0.0%	05_SPCRTU
	04Fall	Priest Rapids Hatchery	OtsPRH13_f a	2013	37	0 – 111	1.5%			0.0%	18_UCOLSF
	04Fall	Priest Rapids Hatchery	OtsPRH14_f a	2014	238	84 – 396	9.6%	265	116 – 446	19.5 %	18_UCOLSF
	04Fall	Priest Rapids Hatchery	OtsPRH15_f a	2015	116	37 – 233	4.7%	149	37 – 260	10.9 %	18_UCOLSF
	04Fall	Lyons Ferry Fish Hatchery	OtsLYON13 S_1_fa	2013			0.0%	82	0 – 201	6.0%	19_SRFALL
	04Fall	Lyons Ferry Fish Hatchery	OtsLYON14 S_1_fa	2014	202	75 – 366	8.2%	85	0 – 169	6.2%	19_SRFALL
	04Fall	Lyons Ferry Fish Hatchery	OtsLYON15 S_1_fa	2015	85	0 – 175	3.4%	85	0 – 210	6.3%	19_SRFALL
	04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH14 S_1_fa	2014			0.0%	42	0 – 126	3.1%	19_SRFALL
	04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH15 S_1_fa	2015			0.0%	38	0 – 115	2.8%	19_SRFALL
	04Fall	Little White Salmon National Fish Hatchery	OtsLWS13_f a	2013	116	37 – 233	4.7%	163	42 – 294	12.0 %	22_BONPOLFA
	04Fall	Little White Salmon National Fish Hatchery	OtsLWS14_f a	2014	126	42 – 252	5.1%	111	0 – 223	8.2%	22_BONPOLFA

	04Fall	Little White Salmon National Fish Hatchery	OtsLWS15_fa	2015	239	83 – 406	9.7%	122	0 – 245	9.0%	22_BONPO OLFA
	04Fall	Umatilla Fish Hatchery	OtsUMA13_fa	2013	37	0 – 111	1.5%			0.0%	23_UMATIL LAFA
	#N/A	#N/A	Unassigned	#N/A	877	598 – 1198	35.5 %			0.0%	#N/A
			TOTAL		2,467		100.0%	1,360		100.0%	

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1167 The fall Non-Treaty commercial fishery is shown by the composition of the combined adult and jack harvest using reporting
 1168 group level (**Table 39**) and broodstock level (**Table 40**) resolution.

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 1170 **Table 39. Summary of the stock composition at the reporting group level of the adult and jack Chinook salmon of the**
 1171 **fall Non-Treaty commercial fishery in 2018 in units of reported catch.**

Reporting Group name	Run type	Reporting Group Code	Hatchery origin- Clipped		Hatchery origin- No Clip		Natural origin- No Clip	
			Estimated abundance		Estimated abundance		Estimated abundance	
			Mean	95% CI	Mean	95% CI	Mean	95% CI
Youngs Bay	Spring	01_YOUNGS						
West Cascade Spring	Spring	02_WCASSP						
West Cascade Fall	Fall	03_WCASFA	190	31 – 647			413	206 – 634
Willamette	Spring	04_WILLAM	30	0 – 124			31	0 – 93
Spring Creek Tule	Fall	05_SPCRTU	3,124	2343 – 3890	146	0 – 387		
Klickitat	Spring	06_KLICKR						
Deschutes spring	Spring	07_DESCSP						
John Day	Spring	08_JOHNDR						
Yakima	Spring	09_YAKIMA						
Upper Columbia spring	Spring	10_UCOLSP						
Tucannon	Spring	11_TUCANO						
Hells Canyon	Spring/Summer	12_HELLSC						
South Fork Salmon	Spring/Summer	13_SFSALM						
Chamberlain Creek	Spring/Summer	14_CHMBLN						
Middle Fork Salmon	Spring/Summer	15_MFSALM						
Upper Salmon	Spring/Summer	16_UPSALM						
Deschutes fall	Fall	17_DESCFA						
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	259	26 – 539	176	0 – 434	2,245	1822 – 2663
Snake River fall	Fall	19_SRFALL	268	32 – 537	391	111 – 744	501	298 – 734
Bonneville Pool spring	Spring	20_BONPOOLSP						

Umatilla spring	Spring	21_UMATILLASP						
Bonneville Pool fall	Fall	22_BONPOOLFA	486	145 – 938	62	0 – 186		
Umatilla fall	Fall	23_UMATILLAFA						
Total			4,356		774		3,190	

Table 40. Summary of the stock composition at the broodstock level of the adult and jack Chinook salmon of the fall Non-Treaty commercial fishery of 2018 in units of reported catch

Fall Commercial Fishery 2018				Adult/jack Chinook AD			Adult/jack Chinook AI			GSI RepGrp
Expected Run Time	Hatchery	Broodstock	Broody ear	ML E	95% CI	Perce nt	ML E	95% CI	Perce nt	
04Fall	Washougal Fish Hatchery	OtsWAS15_fa	2015	141	0 – 424	3.2%			0.0%	03_WCASF A
04Fall	Spring Creek National Fish Hatchery	OtsSPCR15_fa	2015	2,056	1634 – 2483	47.2%	146	32 – 291	18.8%	05_SPCRTU
04Fall	Spring Creek National Fish Hatchery	OtsSPCR16_fa	2016	429	152 – 728	9.9%			0.0%	05_SPCRTU
04Fall	Priest Rapids Hatchery	OtsPRH14_fa	2014	96	0 – 241	2.2%	128	31 – 266	16.5%	18_UCOLSF
04Fall	Priest Rapids Hatchery	OtsPRH15_fa	2015			0.0%	48	0 – 145	6.2%	18_UCOLSF
04Fall	Lyons Ferry Fish Hatchery	OtsLYON13S_1_fa	2013	32	0 – 96	0.7%			0.0%	19_SRFALL
04Fall	Lyons Ferry Fish Hatchery	OtsLYON14S_1_fa	2014	142	31 – 271	3.3%	257	97 – 434	33.2%	19_SRFALL
04Fall	Lyons Ferry Fish Hatchery	OtsLYON15S_1_fa	2015	52	0 – 156	1.2%	85	0 – 189	11.0%	19_SRFALL
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH12S_1_fa	2012			0.0%	49	0 – 146	6.3%	19_SRFALL
04Fall	Little White Salmon National Fish Hatchery	OtsLWS13_fa	2013	176	48 – 338	4.0%	62	0 – 155	8.0%	22_BONPO OLFA
04Fall	Little White Salmon National Fish Hatchery	OtsLWS14_fa	2014	159	48 – 303	3.6%			0.0%	22_BONPO OLFA

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04Fall	Little White Salmon National Fish Hatchery	OtsLWS15_fa	2015	152	0 – 304	3.5%			0.0%	22_BONPO OLFA
		Unassigned		920	515 – 1329	21.1 %			0.0%	
		TOTAL		4,3 56		100.0 %	774		100.0 %	

The fall Treaty Zone 6 commercial fishery is reported by “tule” (i.e. Spring Creek Hatchery tules) and “bright” (upriver bright stocks include 17_DESCFA, 18_UCOLSF, and 19_SRFALL) stocks. The YN monitors sample only the bright stock for DNA analysis and so we provide estimated abundances of the stocks that comprise this Visual Stock Identification “bright” group of fish. It was expected that there would be a small portion of “tule” fish within this group due to some misidentification of the stock using VSI. We estimated 842 tules (05_SPCRTU) which comprised 4% of the total clipped and unclipped hatchery-origin catch. The broodstock composition estimates identified a number of jacks that were included in this harvest from Priest Rapids, Lyons Ferry, and Little White Salmon hatcheries (Broodyear 2016, Table 42)

Table 41. Summary of the stock composition at the reporting group level of the adult VSI-bright Chinook salmon of the fall Treaty zone 6 commercial fishery in 2018 in units of reported catch.

Reporting Group name	Run type	Reporting Group Code	Hatchery origin- Clipped		Hatchery origin- Unclassified	
			Estimated abundance		Estimated abundance	
			Mean	95% CI	Mean	95% CI
Youngs Bay	Spring	01_YOUNGS				
West Cascade Spring	Spring	02_WCASSP				
West Cascade Fall	Fall	03_WCASFA				
Willamette	Spring	04_WILLAM				
Spring Creek Tule	Fall	05_SPCRTU	773	189 – 1612	69	0 – 136
Klickitat	Spring	06_KLICKR				
Deschutes spring	Spring	07_DESCSP				
John Day	Spring	08_JOHNDR				
Yakima	Spring	09_YAKIMA				
Upper Columbia spring	Spring	10_UCOLSP				
Tucannon	Spring	11_TUCANO				
Hells Canyon	Spring/Summer	12_HELLSC				
South Fork Salmon	Spring/Summer	13_SFSALM				
Chamberlain Creek	Spring/Summer	14_CHMBLN				
Middle Fork Salmon	Spring/Summer	15_MFSALM				
Upper Salmon	Spring/Summer	16_UPSALM				
Deschutes fall	Fall	17_DESCFA				
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	6,538	4869 – 8149	1,972	106 – 2839
Snake River fall	Fall	19_SRFALL	3,942	2597 – 5443	2,283	124 – 3442
Bonneville Pool spring	Spring	20_BONPOOLSP				
Umatilla spring	Spring	21_UMATILLASP				
Bonneville Pool fall	Fall	22_BONPOOLFA	3,572	2367 – 4734	516	132 – 860
Umatilla fall	Fall	23_UMATILLAFA	167	0 – 475		
Total			14,992		4,840	

Table 42. Summary of the stock composition at the broodstock level of the adult VSI-bright Chinook salmon of the fall Treaty zone 6 commercial fishery in 2018 in units of reported catch

Fall Treaty Fishery 2018				VSI-Bright Chinook AD			VSI-Bright Chinook AI			GSI RepGrp
Expected Run Time	Hatchery	Broodstock	Brodyear	M L E	95% CI	Percent	M L E	95% CI	Percent	
04Fall	Spring Creek National Fish Hatchery	OtsSPC R15_fa	2015	65 6	262 – 1080	4.4 %	69	0 – 207	1.4 %	05_SPC RTU
04Fall	Priest Rapids Hatchery	OtsPRH 12_fa	2012	18 7	0 – 560	1.2 %			0.0 %	18_UCO LSF
04Fall	Priest Rapids Hatchery	OtsPRH 13_fa	2013	83 3	429 – 1269	5.6 %	12 5	37 – 264	2.6 %	18_UCO LSF
04Fall	Priest Rapids Hatchery	OtsPRH 14_fa	2014	2,0 19	1390 – 2659	13. 5%	1, 24 5	778 – 1783	25. 7%	18_UCO LSF
04Fall	Priest Rapids Hatchery	OtsPRH 15_fa	2015	1,5 17	1024 – 2093	10. 1%	57 2	250 – 962	11. 8%	18_UCO LSF
04Fall	Priest Rapids Hatchery	OtsPRH 16_fa	2016	29	0 – 87	0.2 %	29	0 – 87	0.6 %	18_UCO LSF
04Fall	Lyons Ferry Fish Hatchery	OtsLYO N13S_1_fa	2013	69	0 – 207	0.5 %			0.0 %	19_SRF ALL
04Fall	Lyons Ferry Fish Hatchery	OtsLYO N14S_1_fa	2014	1,9 39	1277 – 2604	12. 9%	1, 26 9	777 – 1802	26. 2%	19_SRF ALL
04Fall	Lyons Ferry Fish Hatchery	OtsLYO N15S_1_fa	2015	97 2	502 – 1470	6.5 %	29 5	101 – 536	6.1 %	19_SRF ALL
04Fall	Lyons Ferry Fish Hatchery	OtsLYO N16S_1_fa	2016	12 2	0 – 365	0.8 %	69	0 – 208	1.4 %	19_SRF ALL
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPF H13S_1_fa	2013	16 6	49 – 333	1.1 %	38 2	135 – 652	7.9 %	19_SRF ALL
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPF H14S_1_fa	2014	29 7	67 – 576	2.0 %	20 1	67 – 401	4.1 %	19_SRF ALL
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPF H15S_1_fa	2015	16 9	0 – 404	1.1 %	67	0 – 201	1.4 %	19_SRF ALL

04Fall	Little White Salmon National Fish Hatchery	OtsLWS 13_fa	2013	44 7	183 – 764	3.0 %	14 9	0 – 373	3.1 %	22_BON POOLF A
04Fall	Little White Salmon National Fish Hatchery	OtsLWS 14_fa	2014	1,9 91	1403 – 2630	13. 3%	31 5	107 – 571	6.5 %	22_BON POOLF A
04Fall	Little White Salmon National Fish Hatchery	OtsLWS 15_fa	2015	1,0 85	692 – 1544	7.2 %	52	0 – 155	1.1 %	22_BON POOLF A
04Fall	Little White Salmon National Fish Hatchery	OtsLWS 16_fa	2016	49	0 – 147	0.3 %			0.0 %	22_BON POOLF A
04Fall	Umatilla Fish Hatchery	OtsUM A13_fa	2013	16 7	44 – 333	1.1 %			0.0 %	23_UM ATILLA FA
#N/A	#N/A	Unassigned	#N/A	2,2 81	1574 – 3052	15. 2%			0.0 %	#N/A
		TOTAL		14, 99 2		10 0.0 %	4, 84 0		10 0.0 %	

The Treaty Sockeye salmon fishery was estimated by clipped hatchery-origin and other unclipped categories (“reintroduced” and natural-origin). We estimated that there were two clipped hatchery stocks, Snake River and Okanogan River stocks (**Table 43**). For the first time, we estimated the number of reintroduced stock from the Yakima River that comprised the fishery (N=149, 2%; **Table 43**). The Yakima River reintroduced stock could be further broken down into broodyear and genetic stock, and was comprised of mostly Wenatchee genetic stock from brood year 2014 (4-year-olds, **Table 44**). The Yakima River sample was small and our confidence intervals overlapped 0. The estimate for these fish at Bonneville Dam was ~1300 fish and so the stock specific harvest rate on this stock would be estimated ~10%. The Treaty harvest rate for the whole run of the sockeye was approximately 4%, and so these numbers seem reasonable given this context

Table 43. Summary of the stock composition at the reporting group level of the sockeye salmon of the Treary fishery in 2018 in units of reported catch

Reporting Group name	Hatchery origin-Clipped		Reintroduction- No Clip		Natural origin- No Clip	
	Estimated abundance		Estimated abundance		Estimated abundance	
	Mean	95% CI	Mean	95% CI	Mean	95% CI
Okanogan	25	0 – 74			6,052	5654 – 6418
Wenatchee					1,312	932 – 1729
SNAKE	150	50 – 299				
Lake Billy Chinook					5	0 – 16

Yakima		149	0 – 558
Odell			34 0 – 103
Total	174	149	7,369

Table 44. Summary of the stock composition at the broodstock level for the reintroduced sockeye salmon from the Yakima River caught in the Treaty fishery in 2018 in units of reported catch

Reintroduced Yakima sockeye					
Stock	Broodyear	Age	MLE	95% CI	Percent
Wenatchee	2014	4	123	0 – 369	82.4%
Okanogan	2013	5	26	0 – 66	17.6%
TOTAL			149		100.0%

Comparison of stock composition among sockeye salmon fisheries

Sockeye salmon were sampled from the lower Columbia River below Bonneville Dam in the lower river sport and above Bonneville Dam in the Zone 6 tribal fishery, and were assigned to five major Columbia River sockeye genetic stocks (Table 45, an additional kokanee stock “Odell” was detected). The lower river commercial harvest did not occur in 2018, and there was no Zone 6 sport fishery in 2018. Low sample numbers of *O. nerka* make it difficult to estimate narrow confidence intervals for abundance estimates of the Yakima River, Snake River, and Lake Billy Chinook stocks (Table 46).

The timing of the sockeye salmon fisheries may influence the harvested proportion of each stock. The Wenatchee stock has an early shifted run in some years like 2018. The Snake River stock (i.e., Redfish Lake) was only represented by 3 fish in the Zone 6 tribal fishery sample (Table 45) making run-timing estimates imprecise for this stock. Of the 3 Snake River fish identified with GSI, all fish were sampled in week 26. Notable difference in stock proportions between Bonneville Dam and the Zone 6 tribal harvest were observed for the Okanogan stock (90% vs. 79%) and for the Wenatchee stock (9% vs. 17%) in the Bonneville Dam vs. harvest mixture samples, respectively (Table 46).

1238 **Table 45. Summary of sample sizes and stock assignments for the 2018 sockeye salmon fisheries by weekly strata.**

Fishery	Stock	Statistical week							
		23	24	25	26	27	28	29	30
Sport	Okanogan					3	3	2	1
Treaty Zone 6	Lake Billy Chinook							1	
	Odell					1			
	Okanogan	3	1	40	33	40	32	25	7
	Snake				3				
	Wenatchee			11	13	4	2	3	1
	Yakima				1			2	

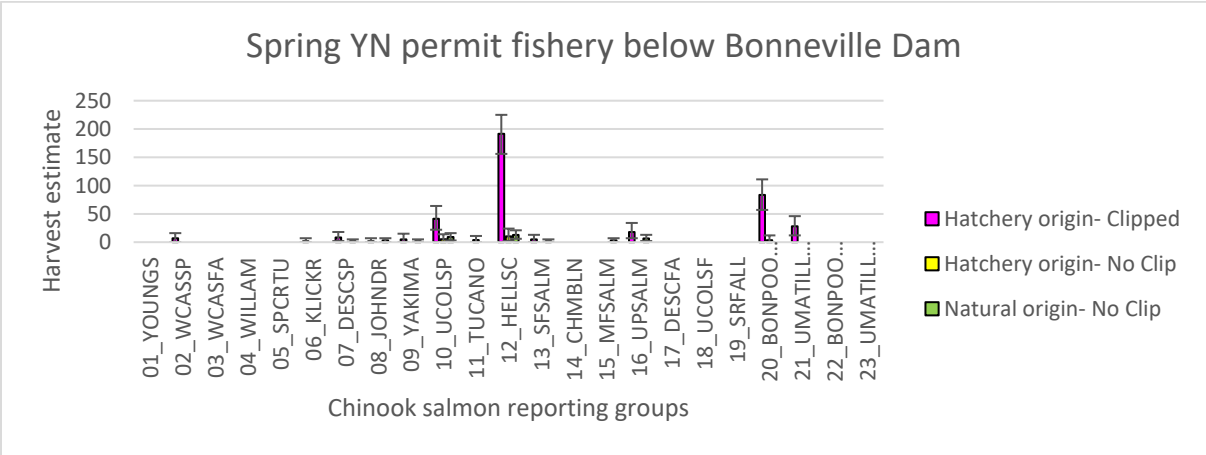
1239

1240 **Table 46. Comparison of stock-specific abundance and percent composition among sockeye salmon fisheries. The mean stock**
1241 **abundance estimate is provided for each fishery harvest in 2018.**

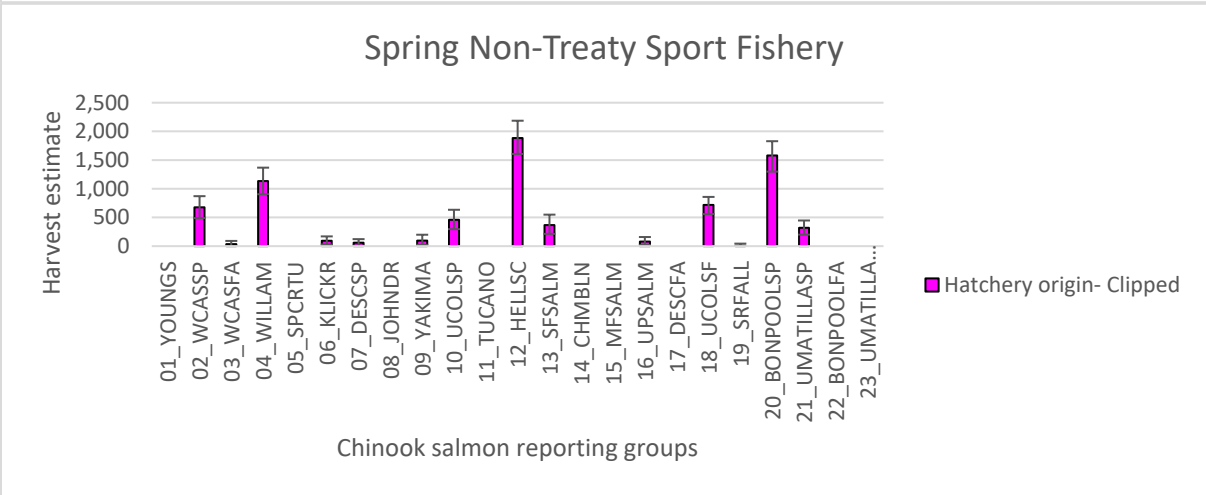
Mixture source	Mean						Stock proportion					
	Okanogan	Wenatchee	Snake	LBC	Yakima	other	Okanogan	Wenatchee	Snake	LBC	Yakima	other
Sport	111	-	-	-	-	-	100.0%	-	-	-	-	-
Treaty Zone 6	6,076	1,312	150	5	149	34	78.6%	17.0%	1.9%	0.1%	1.9%	0.4%
Total Harvest	6,187	1,312	150	5	149	34	78.9%	16.7%	1.9%	0.1%	1.9%	0.4%
Bonneville Dam	174,416	17,675	351	80	1,294	-	90.0%	9.1%	0.2%	0.0%	0.7%	-

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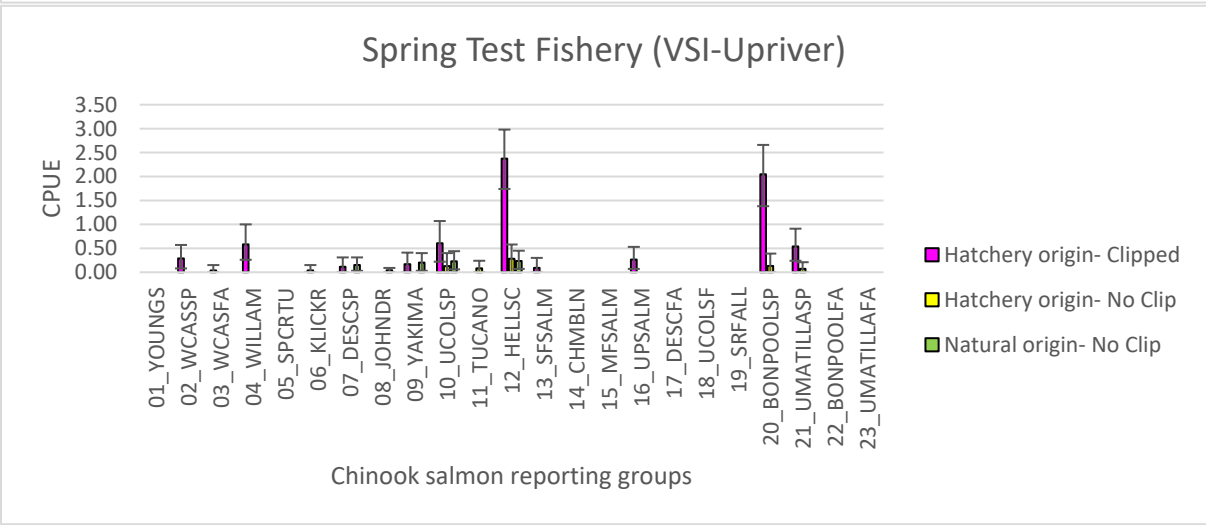
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Figure 21. Stock composition of spring management period clipped and unclipped Chinook salmon harvest mixtures in 2018.

Comparison of stock composition of the clipped and unclipped Chinook salmon stocks among spring-management period Chinook salmon fisheries

The stock composition of the test fishery and the YN permit fishery below Bonneville Dam were very similar (Figure 21) providing further support for the use of the test fishery as a way to predict the upstream stock composition. The sport fishery contained more of the lower river stocks (02_WCASSP and 04_WILLAM) as would be expected given the location of the fishing effort in both regions A and B.

Comparison of stock composition of the clipped and unclipped Chinook salmon stocks between the Treaty and Non-Treaty summer-management period fisheries

Analysis of Chinook salmon fisheries in the summer management period (June 16 – August 1) addressed the following objectives: 1) estimate stock composition for the mark selective sport fishery in Region B and Region A below Bonneville Dam, and 2) compare stock composition of adipose-clipped versus adipose-intact fish from the Tribal Zone 6 harvest above Bonneville Dam. Due to closure of the commercial fishery during the summer management period in 2018, there was no Non-Treaty commercial harvest to compare.

The largest difference in the composition of these two fisheries is the presence of lower river stocks (04_WILLAM) in the lower river fishery compared to the zone 6 fishery (Figure 22). There were small but detectable abundances of Snake River spring stocks in both the Non-Treaty and Treaty summer fisheries.

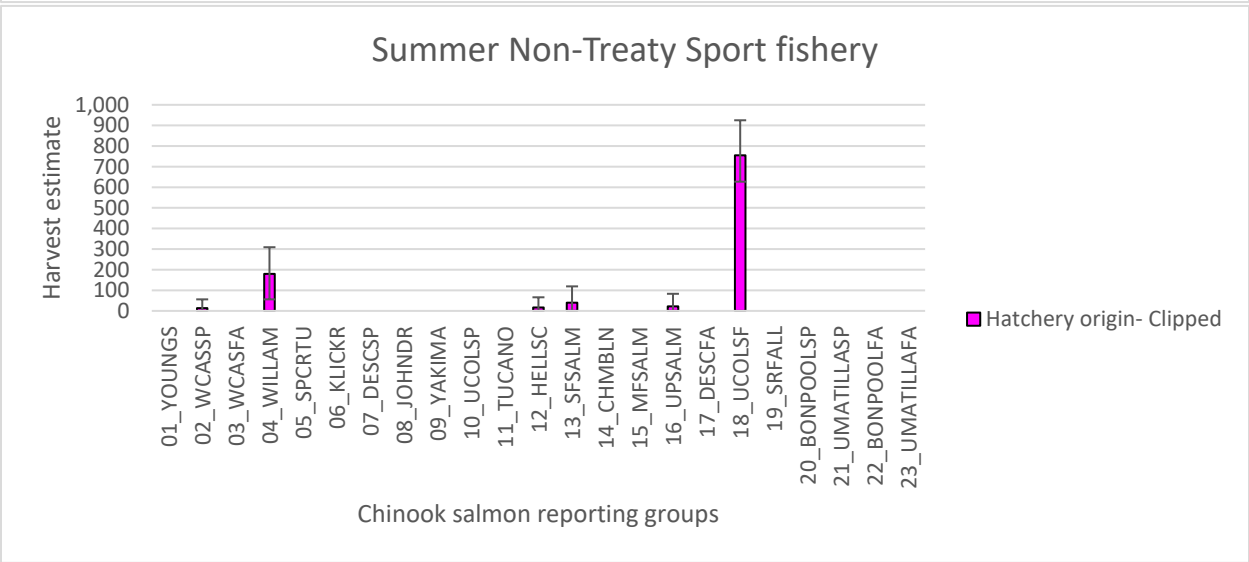
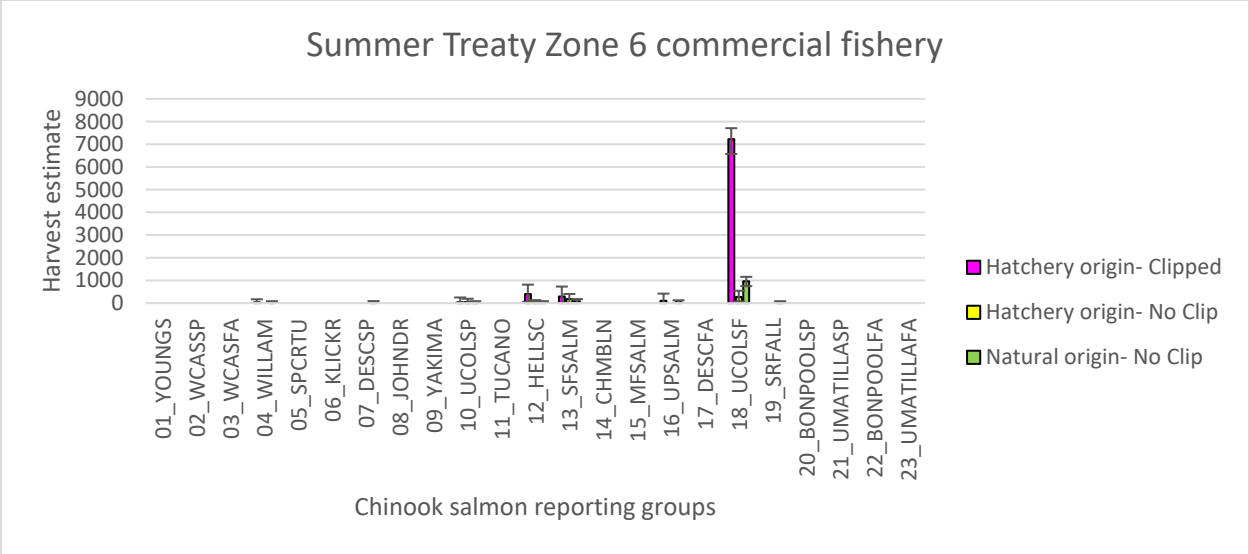


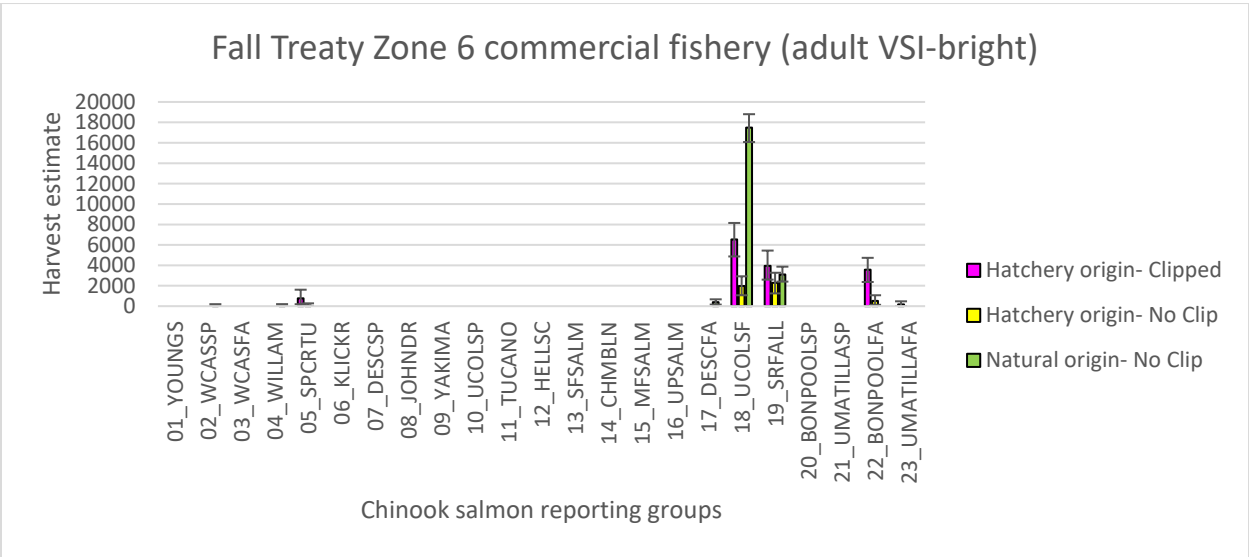
Figure 22. Genetic stock composition of the summer Chinook salmon fisheries analyzed in 2018.

Comparison of stock composition of the clipped and unclipped Chinook salmon stocks among the Treaty and Non-Treaty fall-management period fisheries

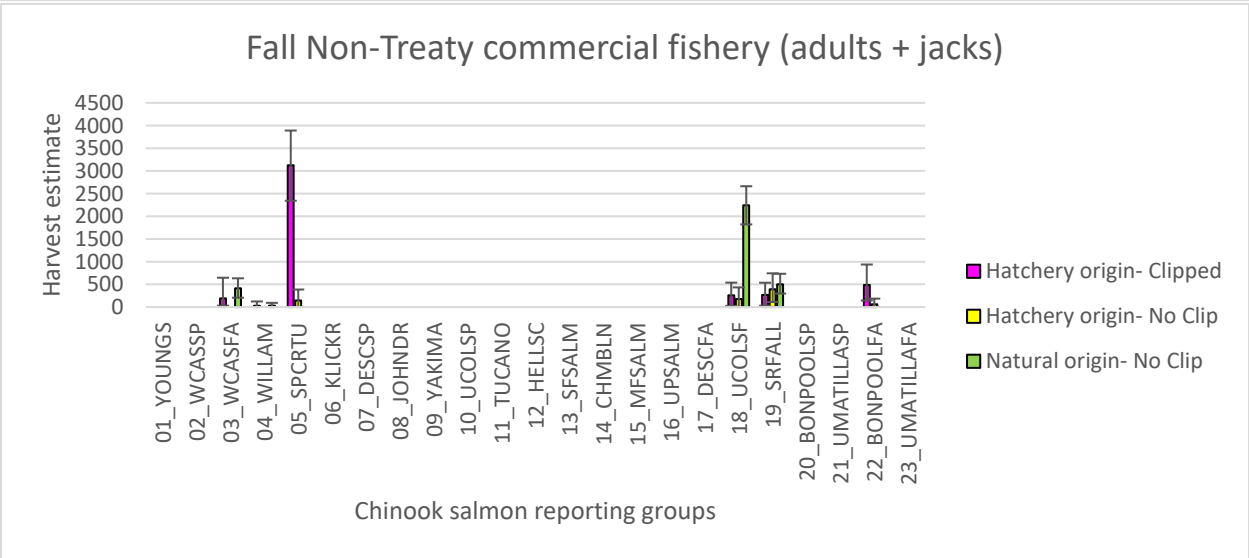
One of the stocks that distinguished the composition of these fall fisheries was the “tule” (05_SPCRTU) stock (Figure 23). There were very low numbers of this stock present in the Treaty Zone 6 fishery due to the way that it was sampled; monitors visually assessed fish as belonging to either “tule” or “bright” category and only DNA sampled the “bright” stock. As such, the Treaty harvest composition can only be used to assess the relative abundance of stocks within the VSI-bright group of fish. However, the fall test fishery (pound net) does provide a more representative composition of the stocks present in the lower river. Comparison of the test fishery composition and the compositions of the commercial and sport harvests showed a

1290 dramatically lower abundance of “tules” in sport fishery compared to the other harvests. This
1291 difference is largely due to sport fishers preferentially keeping the VSI bright fish over the tule
1292 fish.

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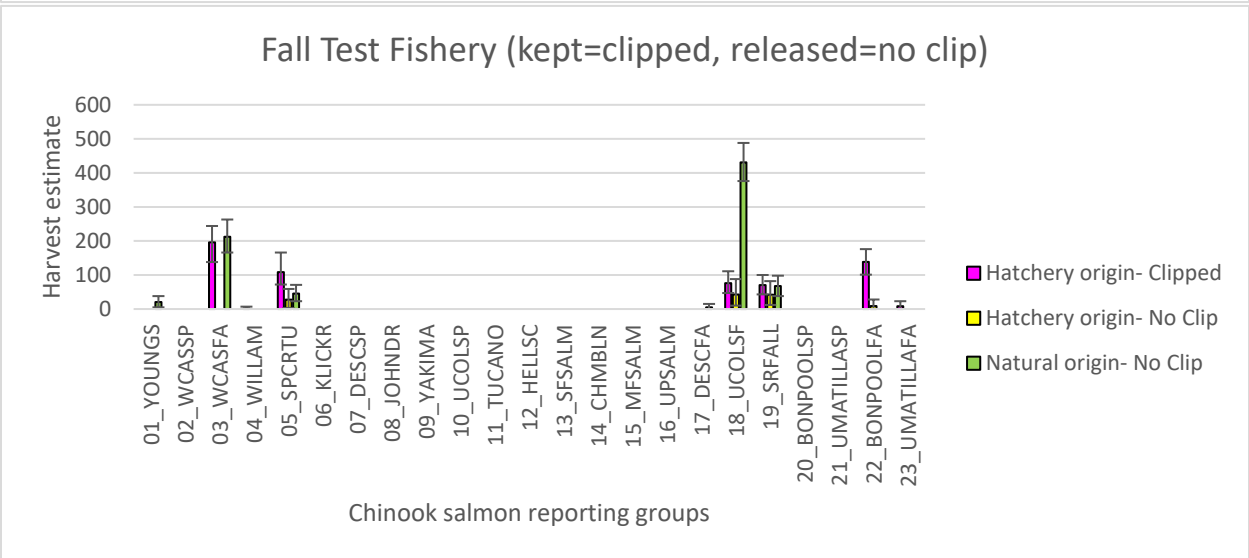
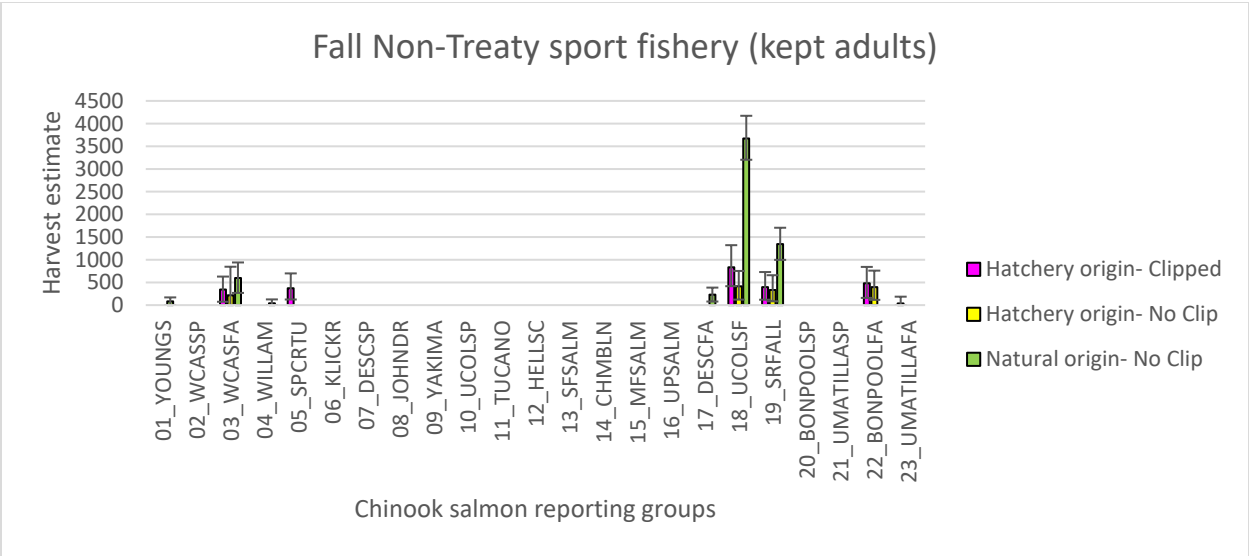


Figure 23. Genetic stock composition of the fall Chinook salmon fisheries analyzed in 2018.

Discussion

Management implications

This study utilized both genetic stock identification (GSI) and parentage based tagging (PBT) in combination to estimate stock composition of mainstem Columbia River Chinook salmon and sockeye fisheries. This is the sixth year in which we were able to assign all three major age classes of spring Chinook from Snake River hatcheries and the second year in which we could assign 3-, 4-, and 5-year old fish to Columbia River hatcheries as a consequence of our expanded PBT baseline. Ongoing expansion of the PBT baseline has now allowed the ability to assign all yearling hatchery jacks and subyearling 4-year-olds originating above Bonneville Dam (complete spawn years date back to 2015), and so future years of analyses will contain all major year classes. Expansion of the PBT baseline to include not only hatcheries of Chinook salmon

and steelhead above Bonneville Dam, but also hatcheries throughout the range of these species could eventually lead to replacing the coded wire tag program for monitoring of in-river harvest stock composition of these species if increases in funding were available and fishery managers thought it were needed. We understand there will be benefit to not only increasing the number of broodstocks represented in the baseline but also expanding the number of SNP loci that we use for PBT applications. A new subset of 254 SNP loci is now being utilized in this report because of its optimal information content across the three major genetic lineages of Chinook Salmon found in the Columbia River Basin and now most broodstocks have complete genotypes. This report includes the sixth genetic analysis on sockeye salmon harvest. This is the first year we have been able to utilize a PBT baseline that can identify offspring of parents from the Yakima River reintroduction. Our results demonstrated differences in stock composition of the sockeye salmon harvest as compared to the total run estimated at Bonneville Dam, but there are questions about the validity of the estimates especially at Bonneville given the potential for sampling error around rare stocks like Snake River sockeye salmon. We will continue to perform PBT and GSI on sockeye salmon harvest in the future to gain further insight into these patterns.

Our most significant achievement in this year, is our ability to expand estimated genetic stock proportions by the reported catch estimates for each fishery. This improvement required tailoring each analysis to the way in which catch is stratified and reported. Our analyses take advantage of new methods (SCOBIDEUX and SPIBETR programs, Delomas and Hess 2020) that are now available to perform tag rate expansions and balance those expansions by proportionally decreasing other stocks in the natural-origin category.

One higher level management question was possible to address in this section:

1) Harvest RM&E: F&W Program Management Question: What are your in-river monitoring results and what are your estimates of stock composition and stock-specific abundance, escapement, catch, and age distribution?

The in-river estimates of stock composition, stock-specific abundance, escapement, catch, and age distribution were addressed for part of the treaty mainstem spring-management period fisheries Chinook salmon harvests below Bonneville Dam, the spring-management test fishery, the non-treaty spring-management sport fishery, the non-treaty and treaty summer-management fisheries, the non-treaty and treaty mainstem fall-management fisheries, and the mainstem treaty sockeye salmon harvests above and non-treaty harvest below Bonneville Dam. For the spring management period of Chinook salmon, we continue to observe a spatial pattern for the stock composition of lower Columbia River stocks which appear more abundant downstream from the Willamette River mouth as compared to upstream of this point which is consistent with a long history of CWT data. We observed differences in the composition of hatchery stocks represented in spring vs. summer management period harvest of Chinook salmon, and run-timing plays an important role in this difference (i.e., late-running stocks appear more abundant among the upriver spring-type lineage that are caught in the summer management period). This pattern is consistent when compared to known origin PIT tagged adult and jack fish tagged as juveniles. Known origin adult age upriver spring and Snake River spring Chinook salmon are almost all past Bonneville by June 15 in most years.

The sockeye salmon tribal fishery is managed in a way that attempts to harvest as many harvestable sockeye salmon as possible under the allowed harvest rate schedule in the U.S. v. Oregon Management Agreement. This 2018 year of analysis of the sockeye salmon harvest

1358 corroborates harvest analyses from previous years, which suggested there may be some over
1359 representation of the Wenatchee sockeye stocks in the Zone 6 harvest as compared to the stock
1360 proportions that are present at Bonneville Dam. The results for Snake River sockeye salmon are
1361 dependent upon representative sampling at Bonneville Dam, but low sample rate and the rarity of
1362 this stock led to uncertainty and high variation around estimates of Snake River sockeye salmon
1363 from Bonneville Dam. Sampling protocols at Bonneville Dam may have higher representation of
1364 young fish as compared to harvest mixtures. Timing of the fishery may also influence the
1365 proportion of each stock, and is consistent with run-timing distributions we observed in previous
1366 reports; the Wenatchee stock has relatively early run-timing but the timing of the Snake River
1367 stock is uncertain due to inconsistent results between PIT-tag and GSI methods. Future analysis
1368 will be needed to examine these patterns for consistency and delve into explanations.
1369 Importantly, we estimated that the Yakima River reintroduction of sockeye has a measurable
1370 (albeit small, ~2%) impact on the Treaty fishery in zone 6.
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Section 4: Characterization of Chinook salmon, sockeye salmon, and steelhead run-timing and abundance at Bonneville Dam

Introduction

The Columbia River Basin supports ESA listed natural-origin stocks of Chinook salmon and steelhead as well as hatchery supplemented populations. Both Chinook salmon and steelhead have been declining in the Columbia River Basin for several reasons including climate change, habitat degradation, hydropower, hatchery practices, and over-harvesting. Along with abundance estimates, basic information related to the way in which stocks of salmonids are spatiotemporally distributed are needed by fisheries managers to achieve sustainable fisheries.

As evident from the genetic stock identification (GSI and PBT) analyses of Chinook and Sockeye salmon fisheries harvests in Section 3, certain stocks seem to have strong spatial and temporal associations. However, because the type of fishing gear, harvest regulations, and the locations targeted varies considerably among fisheries, samples from a representative mixture of all hatchery- and natural-origin stocks at a fixed location is expected to more accurately estimate relative abundance and characterize run-timing distributions of stocks. One potentially ideal fixed location for such sampling is Bonneville Dam, but trapping limitations at this location continue to pose a major challenge for sampling. In addition to information on abundance and run-timing, biological data including fork length and age can be examined with estimated stock of origin to characterize life history differences among stocks. This type of examination is especially important for steelhead, which has been managed using two life-history categories (i.e., A- and B-run). These life-history categories have been observed to be differentially characterized by run-timing at Bonneville Dam (e.g., B-run typically arrives after Aug. 25th), fork length (e.g., by definition, B-run fish are ≥ 78 cm), and ocean age (e.g., most B-run fish tend to spend 2 or more years in saltwater) and all of these types of data have been collected for steelhead in this study.

Project objectives and higher level harvest management questions

Here we analyze fish across the entire run of steelhead, Chinook and Sockeye salmon from April to October to estimate temporally stratified proportions of stocks and extrapolate abundance using a daily census that is conducted at the Bonneville Dam fish counting window. We examine steelhead, Chinook salmon, and Sockeye salmon using sets of species-specific SNP assays for up to 390 loci per species. Although there are some methodological differences among these species-specific applications (e.g., different temporal strata), the general approach to estimating abundance and characterizing run-timing distributions was applied consistently post-season across species in 2018. For all three species, we have demonstrated that these genetic baselines are generally accurate for assigning fish of unknown origin, but the genetic similarity of some stocks requires large reporting groups comprised of broad geographic areas (i.e., mid-Columbia R. and lower Snake R. for spring Chinook salmon). Since Bonneville Dam is the most downstream dam on the Columbia River, the mixture samples obtained here represent the majority of upriver/interior Columbia River Basin stocks. This ongoing study offers a rare opportunity to monitor populations of multiple species of salmonids from a broad geographic range over several years. This long-term study will allow us to characterize trends in run timing and abundance of steelhead, Chinook and sockeye salmon and provide this data to fisheries managers. However, the genetic stock units ('reporting groups') are not the same units that groups of fish are currently managed for due to levels of genetic differentiation that can be detected among baseline stocks (see results under Objective 2 for details). Thus, fisheries

managers continue to explore how to best incorporate genetic monitoring results with more traditional monitoring/tagging programs.

One of the newest features of our analyses is an in-season analysis that was first conducted on Chinook Salmon during the 2017 spring and summer management periods. We have continued offering in-season reports for Chinook Salmon on an approximately bi-weekly report timing schedule during the 2019 spring and summer management periods. Other species and runs have been reported to U.S. v OR Technical Advisory Committee (TAC) members on a timely basis near the end of each management period. Specifically, in 2019, we provided timely post-season reports for steelhead after the Skamania Management Period and the Summer A-/B-Index Management Period, Sockeye Salmon, and Chinook Salmon after the Fall Management Period (see Table 47 for the number and timing of reports for each species and run that were delivered in-season and post-season in 2019).

Harvest RM&E: F&W Program Management Questions:

- i) **What are the status and trend of adult productivity of fish populations?**
- ii) **What are your in-river monitoring results and what are your estimates of stock composition and stock-specific abundance, escapement, catch, and age distribution?**

Analysis of the 2011 dataset by Hess et al. (2012) was the first year we were able to apply Parentage Based Tagging (PBT) to assign a portion of Snake River hatchery-origin spring-run Chinook salmon and summer-run steelhead back to their hatchery parents (Steele et al. 2011). This powerful genetic tool provides the opportunity to obtain additional types of data including accurate age of fish, quantification of the number of non-adipose clipped hatchery-origin fish, and precise assignments of fish to their hatchery broodstock (Steele et al. 2013). The ability of PBT to assign fish to their hatchery broodstock has been shown to be equally accurate as traditional tags (e.g., CWTs; Steele et al. 2013), and PBT provides assignments to specific hatchery broodstocks rather than larger reporting groups used in GSI methods. However, these tools can provide the greatest benefit when applied in combination, as GSI has the ability to provide information on natural-origin fish throughout the Columbia River basin, while PBT is most effective for hatchery-origin fish. The current PBT baseline was recently expanded beyond Snake River hatcheries to include others above Bonneville Dam. However, this effort is ongoing, and while hatcheries continue to be added to our PBT baseline annually, GSI remains a necessary tool for both hatchery- and natural-origin fish that originate from outside the Snake River basin. This report is the 6th year in which all major age classes of steelhead (i.e. 1-, 2-, and 3- ocean ages) and Chinook salmon (3-, 4-, and 5-year olds) can be assigned using the PBT baseline of Snake River hatcheries, and the 4th year in which these can be assigned to some Columbia River hatcheries. This is the first year that Sockeye salmon from the Yakima River reintroduction can be assigned to candidate parents, allowing this stock to be identified in fisheries and at Bonneville Dam. This study integrates PBT and GSI results to provide the greatest amount of stock-specific information available for hatchery- and natural-origin steelhead, Sockeye and Chinook salmon passing Bonneville Dam.

Time line for completion of objectives

Objectives will be ongoing and GSI results updated each year for analyses of salmon and steelhead throughout the accords-funding. This report that is written in January 2020 features the final analyses of 2018, as well as the preliminary in-season and post-season analyses of 2019.

As new genetic techniques are developed they will be applied to this project and results will be compared between years to determine the extent of improvements.

Table 47. The in-season and post-season report timing and scope of the 2019 fish runs.

Species	Management Period	Data coverage	Analysis begins	Report timing
Chinook	Spring	01/01/2019 – 05/03/2019	5/6/2019	5/10/2019
		01/01/2019 – 05/17/2019	5/20/2019	5/24/2019
		01/01/2019 – 06/01/2019	6/3/2019	6/7/2019
		01/01/2019 – 06/15/2019	6/17/2019	6/21/2019
	Summer	01/01/2019 – 07/05/2019	7/8/2019	7/12/2019
		01/01/2019 – 07/31/2019	8/5/2019	8/9/2019
	Fall	01/01/2019 – 10/25/2019	10/28/2019	11/1/2019
Steelhead	Skamania	04/01/2019 – 06/30/2019	7/8/2019	7/12/2019
	Summer A-/B-Index	07/01/2019 – 07/31/2019	8/5/2019	8/9/2019
		07/01/2019 – 10/25/2019	10/28/2019	11/1/2019
Sockeye	Total	01/01/2019 – 08/02/2019	8/5/2019	8/9/2019

Note: The data were reported as cumulative abundance estimates for each genetic stock during the Chinook Salmon and Summer Steelhead A-/B-Index management periods. The report timing indicates the date these reports were provided to the USvOR TAC chair Stuart Ellis for distribution to TAC members.

Methods

Sample Collection

Tissue samples were obtained from adult steelhead (n=878), Chinook (n=3,019) and sockeye salmon (n=1,857) adults in 2018 during migration runs at Bonneville Dam. This sampling effort is covered under the 2008 – 2017 U.S. vs. Oregon harvest biological opinion for sampling at Bonneville Dam.

Sampling for Chinook salmon at Bonneville Dam began during statistical week 17 (04/22/18) and was completed on 10/13/18 (statistical week 41). Sampling occurred at the Adult Fish Facility (AFF) located on the northern end of Bonneville Dam. Fish were sampled 4–5 d per statistical week (except when reduced due to restrictions on trap use or low run size at the beginning and end of the run) and for 4–6 h per day. A picket weir was used to divert migrating fish ascending the Washington shore fish ladder into the AFF collection pool. An attraction flow was used to draw fish through a false weir where they were selected for sampling. After sampling was completed and fish recovered from the anesthetic, they were returned to the Washington shore fish ladder above the picket weir. Only 0.1% of the total Spring management period (i.e., January 1-June 15) adult Chinook salmon count had passed Bonneville by the sampling start date (April 16). In previous years, 2.0-2.5% of the total Spring management period adult Chinook salmon count had passed Bonneville by the sampling start date. However, the adult migration run was delayed passing Bonneville Dam in the spring of 2018 (

Figure 24) and resulted in a lower proportion of unsampled fish. Nonetheless, some early timed stocks may be slightly under-estimated in the results. Restrictions imposed by USACE and NMFS on sampling at the Bonneville AFF result in sample rates for Chinook, sockeye, and

steelhead that are often low. The average sample rate for the adult spring Chinook run in 2018 for the spring and summer management periods was 1.3%, whereas the average sample rate for adult fall Chinook was 0.7% (Table 48).

Based on numbers of fish collected, samples were pooled into weekly strata for Chinook (Table 48), monthly strata for steelhead (Table 49), or a combination thereof for sockeye salmon (Table 50) spanning the majority of the run-year from April to October. We followed a similar protocol as the Monitoring Methods [Protocol "Snake River steelhead and Chinook salmon stock composition estimates \(2010-026-00\) v1.0"](#).

Molecular markers

Expanded panels of genetic markers for steelhead, Sockeye and Chinook salmon are provided in Section 1.

Statistical analyses

Snake River Chinook salmon and steelhead were analyzed for [Parentage assignments using SNPPIT software v1.0](#) (ID: 1341) (Published). The program ONCOR was used to estimate the most likely population-of-origin for the sockeye salmon samples. Individuals were assigned using a ‘best estimate’ approach [Assigning individual samples using Individual Assignment \(IA\) genetic methods v1.0](#) (ID: 1334) (Published). We used GSIsm for [Mixture modeling to estimate stock proportions v1.0](#) (ID: 1333) (Published) to estimate stock composition of Bonneville Dam mixture strata for Chinook salmon and steelhead. Additional detail regarding the specific application to Bonneville Dam are published in Hess et al. (2013).

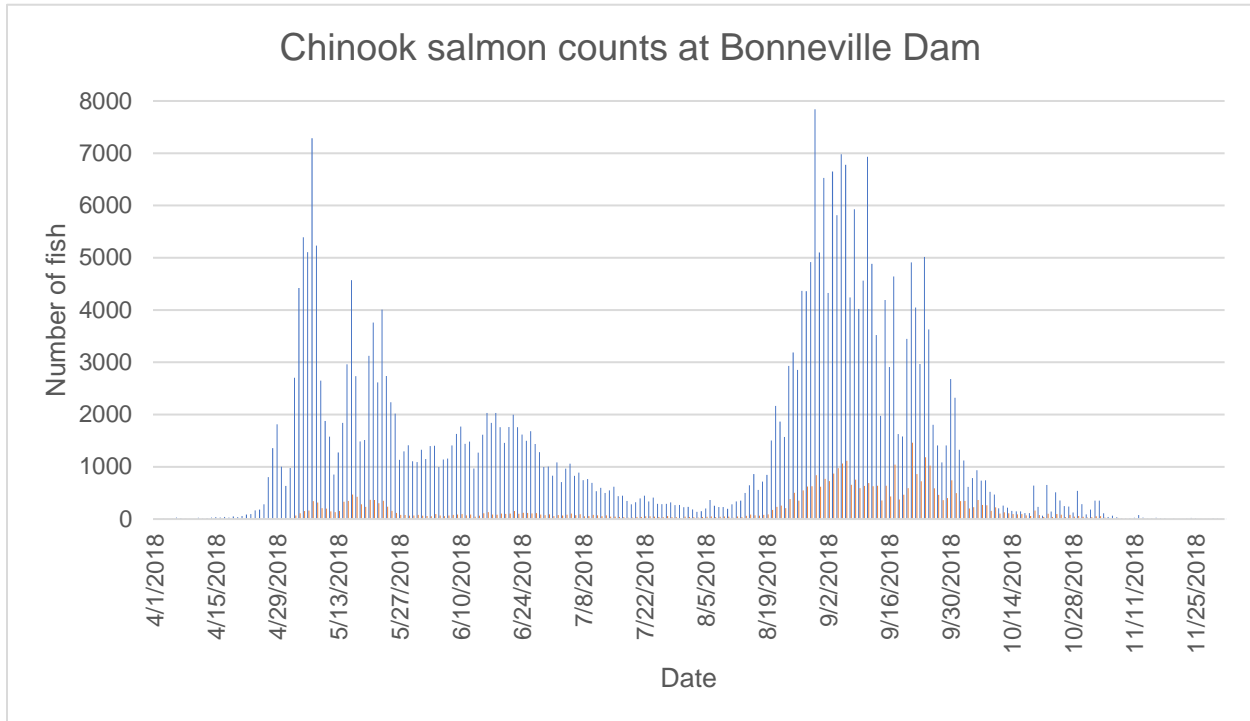


Figure 24. Daily passage of Chinook salmon (adults=blue and jacks=red) at Bonneville Dam in 2018 (source: www.fpc.org).

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Table 48. Sample numbers by weekly strata for Chinook salmon that were DNA sampled or tallied for abundance at Bonneville Dam in 2018.

		Statistical week	TAC		Sample (N)							
			clip count	unclip count	Clipped		Non-clipped		Subtotal		Rate	
					GSI	PBT	GSI	PBT	clip	unclip	clip	unclip
Management period	Spring	1-17	3,088	352	4	52	4	2	56	6	1.8%	1.7%
		18	14,384	2,559	9	142	32	18	151	50	1.0%	2.0%
		19	20,576	4,011	9	186	39	26	195	65	0.9%	1.6%
		20	12,988	3,397	10	158	43	12	168	55	1.3%	1.6%
		21	15,044	5,457	24	181	82	22	205	104	1.4%	1.9%
		22	5,659	2,852	15	64	33	15	79	48	1.4%	1.7%
		23	5,992	3,140	18	81	37	13	99	50	1.7%	1.6%
		24sp	6,058	2,488	13	48	26	7	61	33	1.0%	1.3%
	Summer	24su-25	11,091	3,546	11	45	14	1	56	15	0.5%	0.4%
		26	7,490	2,028	12	38	13	3	50	16	0.7%	0.8%
		27	5,276	1,087	4	17	4	2	21	6	0.4%	0.6%
		28	2,941	1,444	5	37	13	4	42	17	1.4%	1.2%
		29	1,947	909	8	34	19	3	42	22	2.2%	2.4%
		30-31	1,794	1,353	3	25	9	2	28	11	1.6%	0.8%
	Fall	31-34	6,608	13,913	6	18	49	19	24	68	0.4%	0.5%
		35	16,665	19,301	21	71	138	30	92	168	0.6%	0.9%
		36	16,709	24,013	6	32	97	21	38	118	0.2%	0.5%
		37	10,960	19,128	15	52	174	24	67	198	0.6%	1.0%
		38	6,421	16,743	5	34	121	23	39	144	0.6%	0.9%
		39	4,478	12,839	7	28	101	16	35	117	0.8%	0.9%
		40	2,450	7,329	3	20	81	10	23	91	0.9%	1.2%

		41-53	2,226	7,296	1	5	35	5	6	40	0.3%	0.5%
		Total	180,846	155,184	209	1,368	1,164	278	1,577	1,442	0.9%	0.9%

1625 Note: Statistical weeks 1–17 are 1/1/18 – 4/28/18 and 41–53 is 10/7/18–12/31/18. ‘TAC count’ is based on the estimates of clip and unclipped adult
1626 Chinook salmon provided by US v OR Technical Advisory Committee using data from the Fish Passage Center (<http://www.fpc.org>) observed by the
1627 Corps of Engineers at their fish counting window. The sum for each of the clipped and unclipped samples in a given week was used to calculate
1628 sample rate. The management periods approximate the date ranges from January 1st to June 15th (Spring management period), June 16th to July 31st
1629 (Summer management period), and August 1st to December 31 (Fall management period) which are used to categorize spring-, summer-, and fall-run
1630 Chinook salmon, respectively. The number of sampled fish that were assigned via PBT or GSI are shown.

1631 **Table 49. Sample numbers by monthly strata for steelhead that were DNA sampled or tallied for abundance at Bonneville**
1632 **Dam in 2018.**

				Sample (N)											
				A-Index				B-Index				Clipped		Non-clipped	
		Clipped	Non-Clipped	Clipped		Non-clipped		Clipped		Non-clipped		Clipped	Non-clipped	Sample	Sample
	Strata	count	count	GSI	PBT	GSI	PBT	GSI	PBT	GSI	PBT	Total	Total	rate	rate
Skamania	14-26	7,258	5,708	7	43	29	0	0	4	3	0	54	32	0.7%	0.6%
A-/B-Index	27-29	7,515	6,999	17	46	57	2	0	3	1	0	66	60	0.9%	0.9%
	30-34	21,768	12,602	11	69	59	2	0	0	1	0	80	62	0.4%	0.5%
	35-36	14,088	3,684	11	57	13	2	1	32	5	4	101	24	0.7%	0.7%
	37-38	11,626	3,034	1	34	5	2	6	67	7	10	108	24	0.9%	0.8%
	39-40	6,653	1,998	0	26	6	6	9	85	6	20	120	38	1.8%	1.9%
	41-44	2,836	1,197	2	27	8	8	7	39	5	13	75	34	2.6%	2.8%
Summer A-/B-Index subtotal		64,486	29,514	42	259	148	22	23	226	25	47	550	242	0.9%	0.8%
Total		71,744	35,222	49	302	177	22	23	230	28	47	604	274	0.8%	0.8%

1633 Note: Statistical week 14-16 is 4/1/18–6/30/18 (Skamania Management Period) and 27-29 begins the A-/B-Index Period that runs
1634 from 7/1/2018-10/31/2018. ‘Fish count’ is based on tallies of adipose-clipped and unclipped adult steelhead provided by the Fish
1635 Passage Center (<http://www.fpc.org>) observed by the Corps of Engineers at their fish counting window. The total sum of all samples
1636 for each clipped and unclipped category obtained in a given stratum was used to calculate sample rate. The clipped and unclipped
1637 sample numbers were grouped by size (A-Index < 780 mm fork length; B-Index >=780 mm fork length) and further split according to
1638 the number of samples that were either assigned via PBT or GSI.

Table 50. Sample numbers for genetic stock assignments of sockeye salmon that passed Bonneville Dam in 2018.

Statistical week grouping	Bonneville dam fish window count	Genetic stock					Total	Sample rate (%)
		OKA	WEN	RED	LBC	Yakima		
21-23	3290	84	1				85	2.6%
24	17831	214	14				228	1.3%
25	59289	313	37			1	351	0.6%
26	62984	414	42	2		2	460	0.7%
27	34044	277	33				310	0.9%
28	10880	200	20	1			221	2.0%
29	3201	103	7	1	1	1	113	3.5%
30-36	2297	75	10		2	2	89	3.9%
Total	193816	1680	164	4	3	6	1857	1.0%

Note: Statistical week 21 begins on 5/20/18 and 36 ends 9/8/18; there were no sockeye counted before or after these dates. ‘Fish count’ is based on tallies of sockeye salmon adults provided by the Fish Passage Center (<http://www.fpc.org>) observed by the Corps of Engineers at their fish counting window. GSI stocks are Okanagan (OKA), Wenatchee (WEN), Snake River (RED), and Lake Billy Chinook (LBC) and PBT can identify fish from the Yakima reintroduction. The number of samples for a given statistical week or pooled stratum was used to calculate sample rate. Relatively few sockeye salmon were sampled from the RED, LBC, and Yakima stocks, and limits inference regarding run-timing and abundance of these stocks.

Results

Estimated relative abundance of Chinook salmon stocks in 2018

In previous years the 10_UCOLSP reporting group included Carson Hatchery for estimates of relative abundance due to genetic similarity in GSI assignments, so the abundance estimates for this reporting group did not represent actual returns specifically to the upper Columbia River. Beginning in 2017, we have categorized several hatcheries as their own reporting groups to alleviate this issue and so that a more accurate assessment of the number of Chinook returning to the upper Columbia River can be determined. To that end, we have included the following reporting groups that are comprised of collections from our PBT baseline. The 20_BONPOOLSP reporting group includes spring Chinook from Caron Hatchery and Little White Salmon Hatchery. The 21_UMATILLASP reporting group includes spring Chinook from the Umatilla Hatchery. The 22_BONPOOLFA reporting group includes fall Chinook from the Little White Salmon Hatchery. The 22_UMATILLFA reporting group includes fall Chinook from the Umatilla Hatchery.

There were 13 major (i.e., abundance >1000 fish) hatchery origin Chinook salmon stocks represented in the total estimated abundance (N=211,176) of hatchery Chinook salmon passing Bonneville Dam in 2018 (Table 51; Figure 25). These stocks in order of decreasing magnitude were 18_UCOLSF (62,113), 12_HELLSC (35,203), 05_SPCRTU (24,290), 22_BONPOOLFA (22,109), , 19_SRFALL (20,264), , 20_BONPOOLSP (14,028), 16_UPSALM (7,563), 21_UMATILLASP (7,358), 10_UCOLSP (5,592), 13_SFSALM (4,290), 09_YAKIMA (2,943), 02_WCASSP (2,062), and 07_DESCP (1,704) (Table 51; Figure 25). One stocks (13_MFSALM) that was considered major in 2016 did not meet the abundance threshold (>1000) to be considered a major stock in 2017 (Table 51).

With the exception of reporting groups 20_BONPOOLSP, 21_UMATILLASP, 22_BONPOOLFA, and 23_UMATILLAFA), abundance estimates include abundance for PBT-assigned fish (adipose clipped and non-clipped) and adipose clipped fish that were assigned via GSI. PBT assignments improved our ability to accurately identify hatchery origin fish and estimate total stock abundance (Table 51). Further, using PBT assignments we can now provide abundance and run-timing estimates for particular hatchery broodstocks (Table 51) which will allow for much improved abundance estimates.

Table 51. Stock-specific abundance of hatchery origin (adipose clipped and intact) Chinook salmon passing Bonneville Dam in 2018.

			Adult Chinook AD						Adult Chinook AI						GSI RepGrp
Hatchery	Broodstock	Brood year	Spr g	Summe r	Fall	MLE	95% CI	Perce nt	Spr g	Summe r	Fall	MLE	95% CI	Perce nt	
Parkdale Fish Facility	OtsPFF14_sp	2014	2,062	0	0	2,062	1443 – 2732	1.1%	55	0	0	55	0 – 165	0.2%	02_WCASSP
Klickitat Hatchery	OtsKH13_sp	2013	132	0	0	132	0 – 302	0.1%						0.0%	06_KLICKR
Klickitat Hatchery	OtsKH14_sp	2014	722	0	0	722	377 – 1155	0.4%						0.0%	06_KLICKR
Round Butte Fish Hatchery	OtsRB14_sp	2014	1,240	0	0	1,240	728 – 1778	0.7%	52	0	0	52	0 – 157	0.2%	07_DESCSP
Warm Springs National Fish Hatchery	OtsWSNFH14_sp	2014						0.0%	195	0	0	195	0 – 586	0.6%	07_DESCSP
Yakima River Roza Dam	OtsYR14int_sp	2014	2,492	0	0	2,492	1784 – 3252	1.4%						0.0%	09_YAKIMA
Yakima River Roza Dam	OtsYR14seg_sp	2014	451	0	0	451	109 – 851	0.2%						0.0%	09_YAKIMA
Chief Joseph Hatchery	OtsCJH14_sp	2014	477	0	0	477	164 – 828	0.3%	126	0	0	126	0 – 265	0.4%	10_UCOLSP
Eastbank Fish Hatchery	OtsEASTBK14_s p	2014	640	0	0	640	231 – 1080	0.4%	261	0	0	261	0 – 522	0.9%	10_UCOLSP
Eastbank Fish Hatchery	OtsEASTBK15_s p	2015						0.0%	107	0	0	107	0 – 320	0.4%	10_UCOLSP
Leavenworth National Fish Hatchery	OtsLNFH14_sp	2014	1,435	0	0	1,435	820 – 2054	0.8%	73	0	0	73	0 – 219	0.2%	10_UCOLSP
Methow Fish Hatchery	OtsMETH14_sp	2014						0.0%	220	0	0	220	52 – 423	0.7%	10_UCOLSP
Winthrop National Fish Hatchery	OtsWTP13_sp	2013	118	0	0	118	0 – 355	0.1%						0.0%	10_UCOLSP
Winthrop National Fish Hatchery	OtsWTP14_sp	2014	1,574	0	0	1,574	995 – 2161	0.9%	540	0	0	540	289 – 863	1.8%	10_UCOLSP
Lyons Ferry Fish Hatchery	OtsTUCW14S_sp	2014						0.0%	500	0	0	500	231 – 768	1.6%	11_TUCANO
Clearwater Fish Hatchery	OtsPOWP13S_sp	2013	108	0	0	108	0 – 322	0.1%						0.0%	12_HELLSC
Clearwater Fish Hatchery	OtsCLWH14S_sp	2014	7,712	0	0	7,712	6449 – 8974	4.3%	866	43	0	909	547 – 1286	3.0%	12_HELLSC
Clearwater Fish Hatchery	OtsPOWP14S_sp	2014	3,897	0	0	3,897	2961 – 4861	2.2%	434	0	0	434	186 – 685	1.4%	12_HELLSC
Dworshak National Fish Hatchery	OtsDWOR14S_s p	2014	3,841	0	0	3,841	2960 – 4791	2.1%	165	0	0	165	51 – 320	0.5%	12_HELLSC
Lookingglass Fish Hatchery	OtsLOOK14S_sp	2014	540	0	0	540	199 – 913	0.3%	54	0	0	54	0 – 163	0.2%	12_HELLSC
Lookingglass Fish Hatchery	OtsCTHW14S_sp	2014	271	0	0	271	76 – 563	0.2%						0.0%	12_HELLSC
Lookingglass Fish Hatchery	OtsGRUW14S_s p	2014	446	0	0	446	156 – 815	0.2%	302	0	0	302	114 – 542	1.0%	12_HELLSC
Lookingglass Fish Hatchery	OtsLSTW14S_sp	2014	264	0	0	264	82 – 528	0.1%						0.0%	12_HELLSC

Nez Perce	OtsNPFH14S_sp	2014						0.0%	128	0	0	128	0 – 319	0.4%	12_HELLSC
Rapid River Fish Hatchery	OtsRAPH13S_sp	2013	74	0	0	74	0 – 223	0.0%						0.0%	12_HELLSC
Rapid River Fish Hatchery	OtsRAPH14S_sp	2014	15,160	0	0	15,160	13285 – 17001	8.4%	233	0	0	233	75 – 466	0.8%	12_HELLSC
Carson National Fish Hatchery	OtsCAR13_sp	2013	116	0	0	116	0 – 346	0.1%						0.0%	20_BONPOOLS P
Carson National Fish Hatchery	OtsCAR14_sp	2014	4,086	0	0	4,086	3123 – 5122	2.3%	265	0	0	265	67 – 510	0.9%	20_BONPOOLS P
Little White Salmon National Fish Hatchery	OtsLWS13_sp	2013	275	0	0	275	0 – 575	0.2%						0.0%	20_BONPOOLS P
Little White Salmon National Fish Hatchery	OtsLWS14_sp	2014	8,647	226	0	8,873	7408 – 10285	4.9%	412	0	0	412	142 – 695	1.4%	20_BONPOOLS P
Umatilla Fish Hatchery	OtsUMA13_sp	2013	156	0	0	156	0 – 386	0.1%	55	0	0	55	0 – 164	0.2%	21_UMATILLA SP
Umatilla Fish Hatchery	OtsUMA14_sp	2014	6,528	0	0	6,528	5402 – 7684	3.6%	620	0	0	620	332 – 926	2.0%	21_UMATILLA SP
Lookingglass Fish Hatchery	OtsIMNW14S_sp su	2014	628	0	0	628	305 – 1017	0.3%						0.0%	12_HELLSC
McCall Fish Hatchery	OtsMCCA14S_sp su	2014	2,920	394	0	3,314	2372 – 4208	1.8%	696	0	0	696	334 – 1124	2.3%	13_SFSALM
McCall Fish Hatchery	OtsJHNW14S_sp su	2014						0.0%	201	0	0	201	0 – 402	0.7%	13_SFSALM
Pahsimeroi Fish Hatchery	OtsPAHH14S_sp su	2014	1,766	397	0	2,163	1461 – 2911	1.2%						0.0%	16_UPSALM
Sawtooth Fish Hatchery	OtsSAWT14S_sp su	2014	5,058	0	0	5,058	4048 – 6090	2.8%	331	0	0	331	114 – 574	1.1%	16_UPSALM
Chief Joseph Hatchery	OtsCJH13seg_su	2013	292	1,626	0	1,918	1095 – 2781	1.1%	0	330	0	330	96 – 660	1.1%	18_UCOLSF
Chief Joseph Hatchery	OtsCJH13int_su	2013	143	783	0	925	318 – 1606	0.5%						0.0%	18_UCOLSF
Chief Joseph Hatchery	OtsCJH14seg_su	2014	1,424	6,713	0	8,137	5520 – 9726	4.5%						0.0%	18_UCOLSF
Chief Joseph Hatchery	OtsCJH14int_su	2014	574	3,902	0	4,476	3277 – 5724	2.5%	141	548	0	689	213 – 1237	2.3%	18_UCOLSF
Chief Joseph Hatchery	OtsCJH15int_su	2015	0	0	489	489	0 – 1466	0.3%						0.0%	18_UCOLSF
Eastbank Fish Hatchery	OtsEASTBK12_s u	2012	312	261	0	573	104 – 1136	0.3%						0.0%	18_UCOLSF
Eastbank Fish Hatchery	OtsEASTBK13_s u	2013	614	1,411	0	2,025	1195 – 2928	1.1%	73	246	0	318	73 – 638	1.0%	18_UCOLSF
Eastbank Fish Hatchery	OtsEASTBK14_s u	2014	1,594	3,749	194	5,537	4173 – 6796	3.1%	148	135	0	283	81 – 533	0.9%	18_UCOLSF
Entiat National Fish Hatchery	OtsENFH13_su	2013	0	164	0	164	0 – 486	0.1%	0	141	0	141	0 – 424	0.5%	18_UCOLSF
Entiat National Fish Hatchery	OtsENFH14_su	2014	737	5,768	0	6,505	4667 – 8078	3.6%	174	0	0	174	0 – 411	0.6%	18_UCOLSF
Wells Fish Hatchery	OtsWELLS12_su	2012						0.0%	63	0	0	63	0 – 190	0.2%	18_UCOLSF
Wells Fish Hatchery	OtsWELLS13_su	2013	1,646	274	0	1,919	1300 – 2646	1.1%	77	0	0	77	0 – 232	0.3%	18_UCOLSF
Wells Fish Hatchery	OtsWELLS14_su	2014	1,155	4,581	0	5,736	4376 – 7136	3.2%	77	243	0	320	0 – 806	1.1%	18_UCOLSF

Wells Fish Hatchery	OtsWELLS15_su	2015	63	154	0	217	0 – 524	0.1%						0.0%	18_UCOLSF
Spring Creek National Fish Hatchery	OtsSPCR15_fa	2015	0	0	15,555	15,555	12630 – 18525	8.6%	0	0	867	867	237 – 1589	2.9%	05_SPCRTU
Spring Creek National Fish Hatchery	OtsSPCR16_fa	2016	0	0	2,989	2,989	1345 – 4999	1.7%	0	0	321	321	0 – 962	1.1%	05_SPCRTU
Priest Rapids Hatchery	OtsPRH13_fa	2013	0	0	1,145	1,145	487 – 1822	0.6%	0	0	995	995	405 – 1696	3.3%	18_UCOLSF
Priest Rapids Hatchery	OtsPRH14_fa	2014	0	0	2,979	2,979	1674 – 4294	1.6%	0	0	3,079	3,079	2136 – 4310	10.2%	18_UCOLSF
Priest Rapids Hatchery	OtsPRH15_fa	2015	0	0	3,900	3,900	2604 – 5323	2.2%	0	0	3,602	3,602	2421 – 4813	11.9%	18_UCOLSF
Lyons Ferry Fish Hatchery	OtsLYON13S_1_fa	2013	0	0	844	844	187 – 1501	0.5%	0	0	547	547	118 – 1027	1.8%	19_SRFALL
Lyons Ferry Fish Hatchery	OtsLYON14S_1_fa	2014	0	64	3,452	3,516	2201 – 5055	1.9%	0	165	3,289	3,455	2360 – 4636	11.4%	19_SRFALL
Lyons Ferry Fish Hatchery	OtsLYON15S_1_fa	2015	0	0	5,085	5,085	3121 – 7089	2.8%	0	0	1,580	1,580	890 – 2357	5.2%	19_SRFALL
Lyons Ferry Fish Hatchery	OtsLYON16S_1_fa	2016						0.0%	0	0	217	217	0 – 651	0.7%	19_SRFALL
Nez Perce Tribal Fish Hatchery	OtsNPFH13S_1_fa	2013						0.0%	0	0	205	205	0 – 615	0.7%	19_SRFALL
Nez Perce Tribal Fish Hatchery	OtsNPFH14S_1_fa	2014	0	0	1,172	1,172	440 – 2092	0.6%	0	0	1,248	1,248	525 – 2088	4.1%	19_SRFALL
Nez Perce Tribal Fish Hatchery	OtsNPFH15S_1_fa	2015	0	65	170	235	0 – 575	0.1%	0	0	1,602	1,602	770 – 2472	5.3%	19_SRFALL
Little White Salmon National Fish Hatchery	OtsLWS13_fa	2013	0	0	4,848	4,848	2925 – 6881	2.7%	0	0	1,311	1,311	589 – 2087	4.3%	22_BONPOOLFA
Little White Salmon National Fish Hatchery	OtsLWS14_fa	2014	0	0	6,839	6,839	5093 – 8681	3.8%	0	0	1,628	1,628	942 – 2434	5.4%	22_BONPOOLFA
Little White Salmon National Fish Hatchery	OtsLWS15_fa	2015	0	0	7,144	7,144	5375 – 8960	4.0%	0	0	339	339	102 – 685	1.1%	22_BONPOOLFA
#N/A	Unassigned	#N/A	1,403	8	9,711	11,122	8963 – 15354	6.1%						0.0%	#N/A
	TOTAL		83,790	30,539	66,516	180,846		100.0%	7,647	1,852	20,831	30,330		100.0%	

1681 **Note:** These summary statistics of run-timing distributions were calculated using a method to estimate abundance of each stock based on temporally stratified stock proportions and TAC
1682 estimates of clipped and unclipped Chinook salmon that passed Bonneville Dam at the fish counting window. Abundance was reported for each stock as subtotals that passed within each
1683 management period.

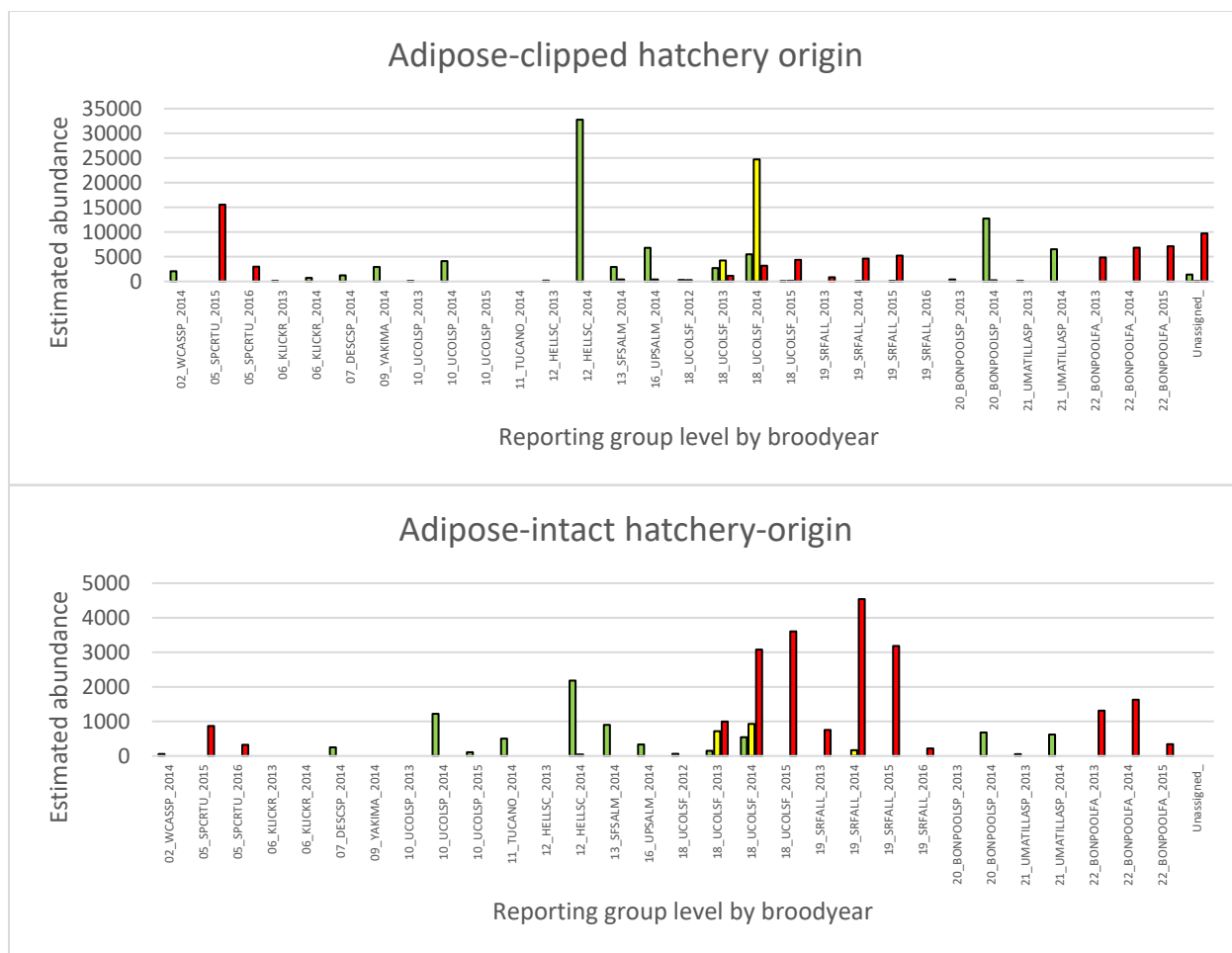


Figure 25. Estimated abundance of hatchery origin adipose-clipped (top) and adipose-intact (bottom) Chinook assigned by PBT reporting groups and broodyears that were sampled at Bonneville Dam in 2018 during spring (green), summer (yellow) and fall (red) management periods.

In Figure 25, there are several new reporting groups that have been created and are populated based on hatchery fish assigned via PBT, which were subsequently categorized into higher level groups to complement the GSI reporting groups. For example, 20_BONPOOLSP reporting group includes PBT assignments of spring Chinook to Carson Hatchery and Little White Salmon Hatchery. The 21_UMATILLASP reporting group includes PBT assignments of spring Chinook to the Umatilla Hatchery. The 22_BONPOOLFA reporting group includes PBT assignments of fall Chinook to the Little White Salmon Hatchery. The 22_UMATILLFA reporting group includes PBT assignments of fall Chinook to the Umatilla Hatchery. These reporting groups serve the purpose of distinguishing these hatchery broodstocks apart from other reporting groups that share genetic affinity with these hatcheries but are part of ESA listed groups. For example, 20_BONPOOLSP broodstocks share genetic similarity with upper Columbia River spring Chinook Salmon, but only hatchery fish from the latter group are ESA listed.

We detected PBT assignments for 19.3% (278/1442) of adipose non-clipped (i.e., presumed natural-origin) Chinook salmon sampled at Bonneville Dam in 2017. There were 7

1706 major (i.e., abundance >1000 fish) Chinook salmon stocks represented in the total estimated
1707 relative abundance (N=124,854) of natural origin (i.e., adipose non-clipped fish that did not
1708 assign via PBT) Chinook salmon passing Bonneville Dam in 2018 (Table 52; Figure 26). These
1709 non-clipped stocks in order of decreasing magnitude were 18_UCOLSF (92,425), 19_SRFALL
1710 (15,406), 12_HELLSC (4,833), 10_UCOLSP (3,607), 09_YAKIMA (2,240), 17_DESCFA
1711 (1,752), and 13_SFSALM (1,402). These stock abundance estimates were based on the stock
1712 proportions that were estimated in using SCOBIDEUX and SPIBETR functions and the
1713 estimates of clipped and unclipped adults distributed by TAC (Table 48).

1714 **Table 52. Estimated abundance of natural origin Chinook salmon stocks passing**
1715 **Bonneville Dam in 2018.**

						Natural origin- No Clip	
Reporting Group name	Run type	Reporting Group Code	Spring	Summer	Fall	Estimated abundance	
						Mean	95% CI
Youngs Bay	Spring	01_YOUNGS	0	85	97	182	0 – 448
West Cascade Spring	Spring	02_WCASSP					
West Cascade Fall	Fall	03_WCASFA					
Willamette	Spring	04_WILLAM	0	0	204	204	0 – 613
Spring Creek Tule	Fall	05_SPCRTU	0	0	526	526	160 – 1071
Klickitat	Spring	06_KLICKR	52	0	0	52	0 – 157
Deschutes spring	Spring	07_DESCSP	745	0	0	745	427 – 1060
John Day	Spring	08_JOHNDR	51	0	0	51	0 – 154
Yakima	Spring	09_YAKIMA	2,240	0	0	2,240	1685 – 2815
Upper Columbia spring	Spring	10_UCOLSP	3,607	0	0	3,607	2862 – 4306
Tucannon	Spring	11_TUCANO	207	0	0	207	51 – 400
Hells Canyon	Spring/Summer	12_HELLSC	4,833	0	0	4,833	4104 – 5611
South Fork Salmon	Spring/Summer	13_SFSALM	748	654	0	1,402	779 – 2103
Chamberlain Creek	Spring/Summer	14_CHMBLN					
Middle Fork Salmon	Spring/Summer	15_MFSALM	525	0	0	525	278 – 827
Upper Salmon	Spring/Summer	16_UPSALM	655	41	0	696	385 – 1025
Deschutes fall	Fall	17_DESCFA	0	0	1,752	1,752	1060 – 2478
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	2,885	6,950	82,590	92,425	88839 – 95299
SNAKE RIVER fall	Fall	19_SRFALL	59	785	14,562	15,406	13193 – 17744
Total			16,608	8,515	99,731	124,854	

1716 Note: Abundance of natural-origin stocks is shown by subtotals that passed Bonneville Dam in
1717 each management period (spring, summer, and fall) and the total mean estimate with 95% C.I.
1718 Natural-origin estimates exclude abundance of adipose-intact hatchery-origin fish.

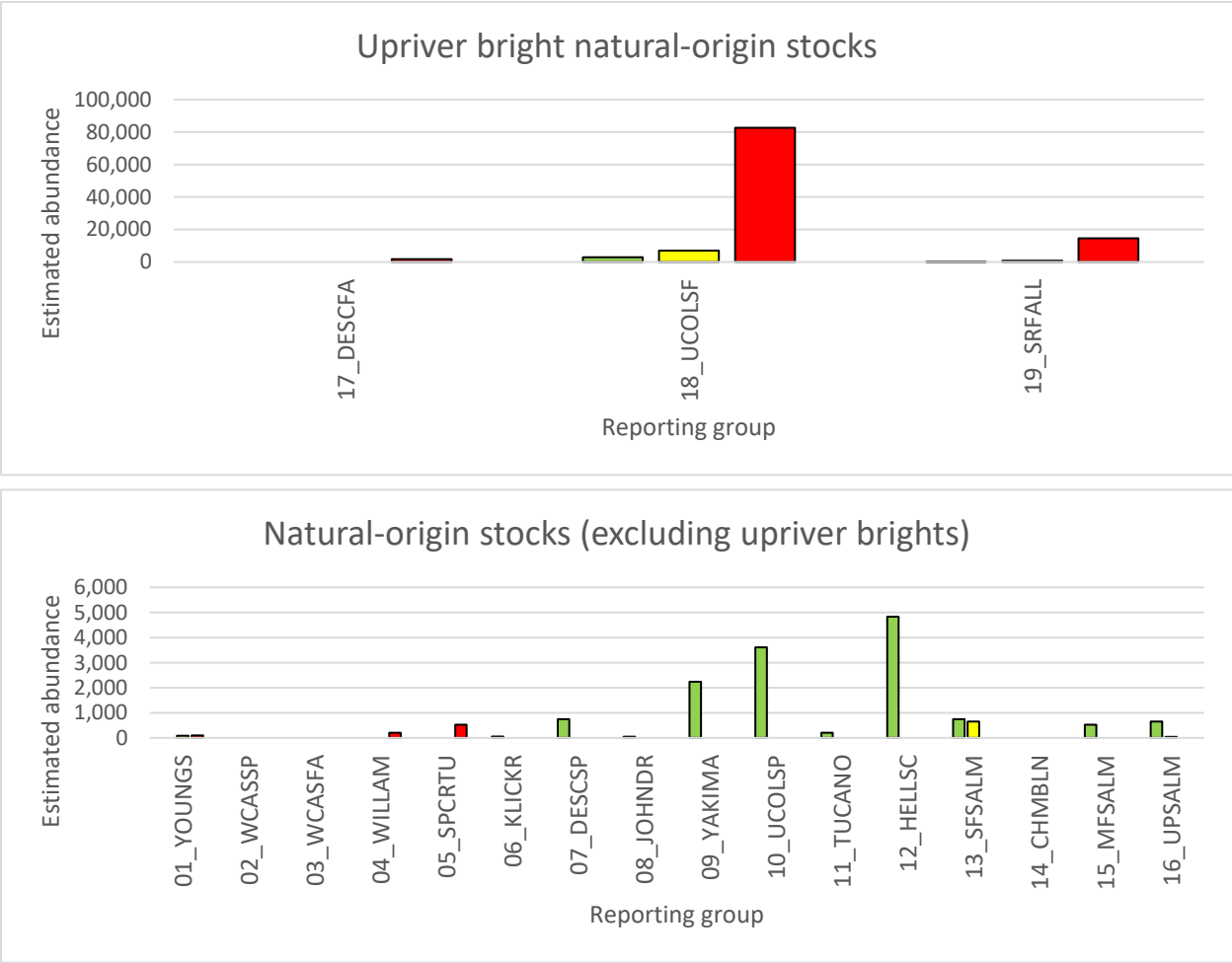
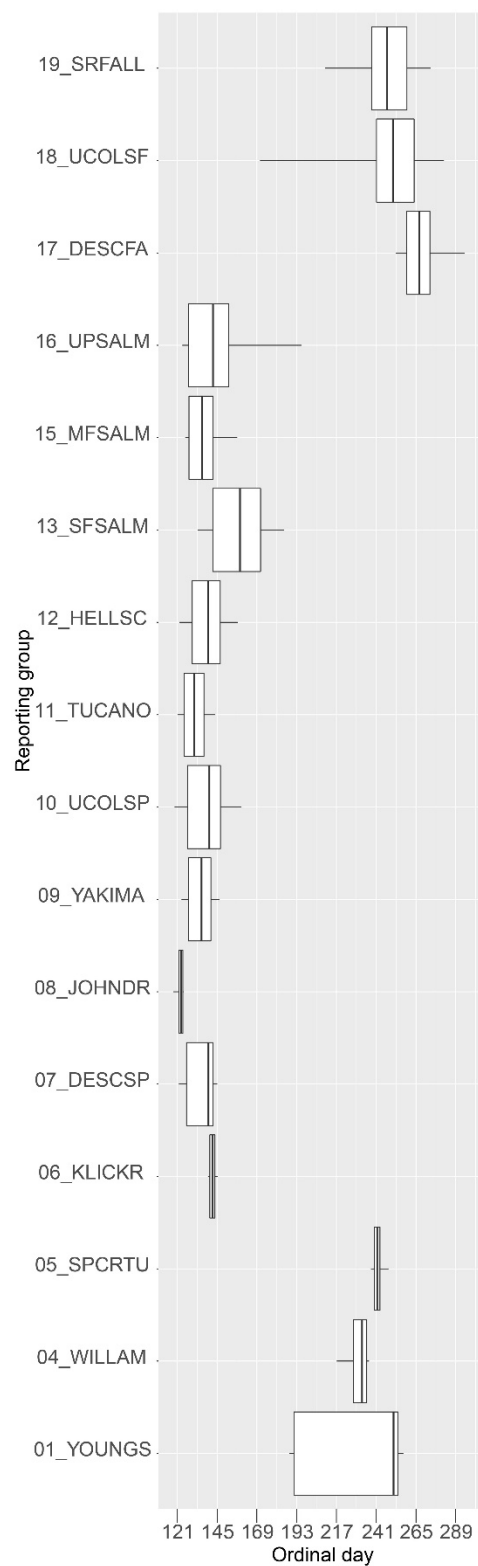


Figure 26. Estimated abundance of natural origin (excluding adipose-intact hatchery-origin fish) Chinook sampled at Bonneville Dam in 2018 assigned to genetic stock of origin. Upriver bright Chinook salmon reporting groups (top panel), and all other natural-origin Chinook reporting groups (bottom panel) are shown by spring (green), summer (yellow) and fall (red) management period.

Run-timing of Chinook salmon stocks in 2018

We plotted the run-timing distributions of the natural-origin (excluding adipose unclipped hatchery-origin fish) Chinook salmon stocks and provide subtotals of abundance for each management period (Table 52; **Figure 27**). While the median date of passage for all natural-origin spring Chinook stocks occurred well within the spring management period, the run-timing for 13_SFSALM and 16_UPSALM was found to extend beyond the spring management period (i.e., the 95th percentile of their run distribution occurred after 6/16/17; ordinal day 166). We estimated that 47% and 6% of the total abundance of these two spring stocks arrived during the summer management period for 3_SFSALM and 16_UPSALM, respectively (Table 52). The run-timing for one natural-origin summer/fall Chinook stock (i.e., 18_UCOLSF) and three natural-origin fall Chinook salmon stocks (i.e., 05_SPCRTU, 17_DESCFA, and 19_SRFALL) all had median dates on or after ordinal day 241 (8/29/18, **Figure 27**). We estimated that for the 18_UCOLSF stock abundance that arrives during the spring and summer management periods, approximately 29% arrives during the spring (Table 52).

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Figure 27. Run-timing distributions (median ordinal day, interquartile range, and 5th and 95th percentile) for natural origin Chinook (excluding all adipose-intact hatchery-origin fish) assigned to reporting group of origin that were sampled at Bonneville Dam in 2018.

Estimated relative abundance of steelhead stocks in 2018

Daily passage of steelhead at Bonneville Dam in 2018 is provided in Figure 28. There were five major stocks (abundance >1000) represented in the total estimated relative abundance (N=70,265) of hatchery origin steelhead passing Bonneville Dam in 2018 (Table 53). These stocks in order of decreasing magnitude were 10_SFCLWR (26,807), 07_MGILCS (18,878), 14_UPSALM (17,707), 09_UPPCOL (4,637), and 03_SKAMAN (2,101) (Table 53; Figure 29). These same stocks were identified as being of major abundance in 2016 and 2017. These estimates include relative abundance estimated from PBT-assigned fish that were mostly adipose clipped; however, a portion of the PBT-assigned fish were found to be non-clipped. Therefore, PBT assignments improved our ability to accurately identify hatchery-origin steelhead and estimate total stock relative abundance. Further, using PBT assignments we can now provide abundance (Table 54; Figure 30) and run-timing estimates for particular hatchery broodstocks (Table 54). There were 12 major hatchery broodstock sources (abundance >1000) represented in the total estimated abundance of hatchery-origin steelhead passing Bonneville Dam in 2018 (Table 54). These stocks in order of decreasing magnitude were OmyDWOR15S (25,501), OmyLYON16S (5,657), OmyWALL15S (4,461), OmyOXBO16S (4,395), OmyPAHH16S (3,533), OmyOXBO15S (3,433), OmyLYON15S (2,822), OmySAWT16S (2,784), OmyWEL_OKA15 (2,516), OmyPAHH15S (2,288), OmySKH14_su (1,386), and OmyRB15 (1,167).

We detected PBT assignments for 28.5% (69/242) of adipose non-clipped (i.e., presumed natural-origin) steelhead sampled at Bonneville Dam in 2018. There were three major stocks (abundance >1000) represented in the total estimated relative abundance (N=23,735) of natural origin (excluding adipose unclipped hatchery-origin fish) steelhead passing Bonneville Dam in 2018 (Table 53). These stocks in order of decreasing magnitude were 07_MGILCS (16,797), 09_UPPCOL (1,424), and 08_YAKIMA (1,321). Two major stocks in 2017 were no longer estimated with abundance >1000: 14_UPSALM (35, down from 2,286), and 06_KLICKR (942, down from 1,013). The 14_UPSALM stock appeared to have decreased dramatically but we feel this was a results of the SCOBIDEUX SPIBETR function that minimized bias from tag rate expansion of the unclipped hatchery-origin fish. This results underscores the importance of the SPIBETR method and using this function consistently on past years of data for better comparisons across years.

1776 **Table 53. Stock-specific abundance by A-/B-Index categories of hatchery- (adipose clipped and non-clipped) and natural-origin summer A-/B-Index steelhead passing**
1777 **Bonneville Dam in 2018.**

Reporting Group name	Reporting Group Code	Hatchery origin- Clipped				Hatchery origin- No Clip				Natural origin- No Clip			
		A-INDEX		B-INDEX		A-INDEX		B-INDEX		A-INDEX		B-INDEX	
		Mean	95% CI	Mean	95% CI	Mean	95% CI	Mean	95% CI	Mean	95% CI	Mean	95% CI
Lower Columbia	02_LOWCOL	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0
Skamania	03_SKAMAN	1,995	572 – 3876	107	0 – 449	0	0 – 0	0	0 – 0	467	117 – 817	0	0 – 0
Willamette	04_WILLAM	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0
Big White Salmon	05_BWSALM	139	0 – 558	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0
Klickitat	06_KLICKR	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	792	266 – 1396	151	0 – 371
mid Columbia/Grande Ronde/Imnaha/lower Clearwater/lower Snake/lower Salmon	07_MGILCS	18,199	14618 – 22230	58	0 – 233	621	36 – 1637	0	0 – 0	16,760	15310 – 18228	36	0 – 117
Yakima	08_YAKIMA	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	1,321	560 – 2073	0	0 – 0
upper Columbia	09_UPPCOL	4,372	2254 – 6883	0	0 – 0	265	0 – 796	0	0 – 0	1,067	407 – 1780	357	0 – 763
SF Clearwater	10_SFCLWR	3,253	1897 – 4809	19,159	16873 – 21374	576	139 – 1117	3,819	2546 – 5181	386	62 – 851	370	60 – 913
upper Clearwater	11_UPCLWR	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	248	51 – 498	398	107 – 716
SF Salmon	12_SFSALM	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	467	105 – 894
MF Salmon	13_MFSALM	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	627	153 – 1187	253	0 – 506
upper Salmon	14_UPSALM	16,773	12692 – 20773	430	0 – 1062	498	0 – 1215	0	0 – 0	35	0 – 104	0	0 – 0
	Total	44,732		19,754		1,960		3,819		21,703		2,032	

1778 **Note:** These summary statistics of run-timing distributions were calculated using a method to estimate abundance of each stock based on temporally stratified stock proportions
1779 and TAC estimates of clipped and unclipped steelhead that passed Bonneville Dam at the fish counting window. This method for estimating abundance minimizes bias imposed
1780 by uneven sampling.

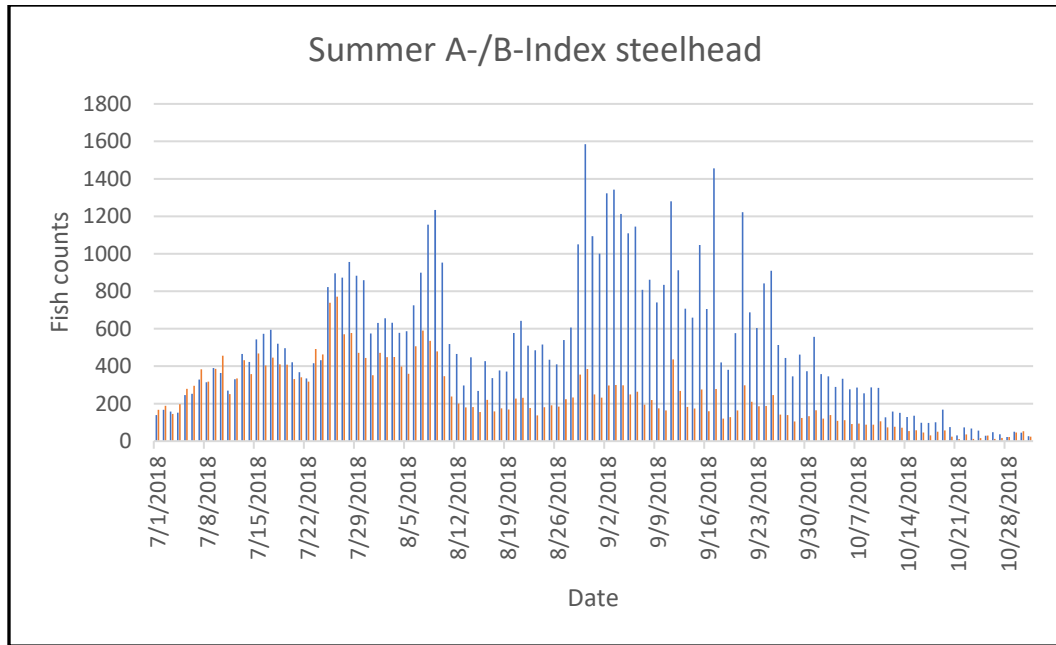
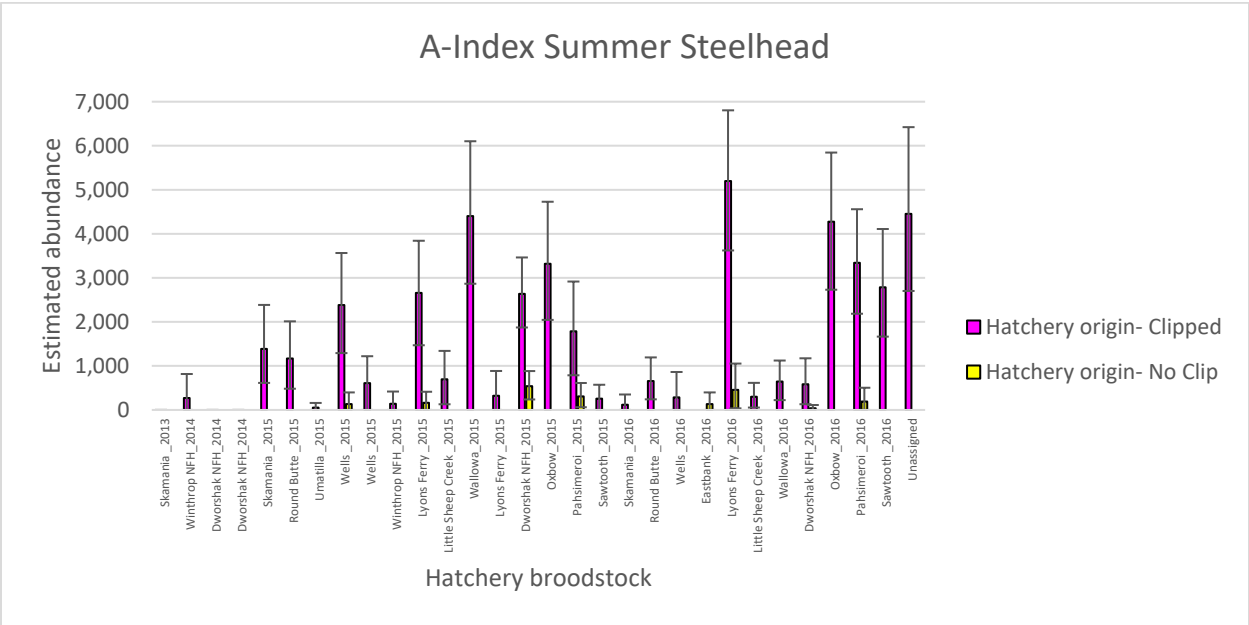


Figure 28. Daily passage of clipped (blue) and unclipped (red) steelhead at Bonneville Dam in 2018 during the summer A-/B-Index management period (source: www.fpc.org).



Figure 29. Estimated abundance (\pm 95% CI) of A-Index (<780mm FL, top) and B-Index (\geq 780mm FL, bottom) hatchery origin (adipose clipped and unclipped) and natural-origin steelhead assigned to genetic stock of origin that were sampled at Bonneville Dam in 2018.

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Figure 30. Estimated abundance (\pm 95% CI) of A-Index (<780mm FL, top) and B-Index (\geq 780mm FL, bottom) hatchery origin steelhead (adipose clipped and unclipped) sampled at Bonneville Dam in 2018 that assigned via PBT to 29 hatchery broodstocks of origin. The 2016 age-class (1-ocean fish), 2015 age-class (2-ocean fish), and 2014 age class (3-ocean fish) are shown. Key to broodstock collection is presented in Appendix 7.

1798 **Table 54. Hatchery broodstock-specific estimated abundance of A-Index and B-Index adipose clipped and non-clipped PBT-assigned steelhead passing Bonneville Dam in 2018.**

					Hatchery clipped				Hatchery Unclipped			
					A-INDEX		B-INDEX		A-INDEX		B-INDEX	
Hatchery	Stock	GSI RepGrp	Broodstock	Broodyear	MLE	95% CI	MLE	95% CI	MLE	95% CI	MLE	95% CI
Skamania	Summer	03_SKAMAN	OmySKH12_su	2013	0	0 – 0	107	0 – 326				
Skamania	Summer	03_SKAMAN	OmySKH14_su	2015	1,386	614 – 2385	0	0 – 0				
Skamania	Summer	03_SKAMAN	OmySKH15_su	2016	117	0 – 351	0	0 – 0				
Round Butte	Deschutes River	07_MGILCS	OmyRB15	2015	1,167	482 – 2012	0	0 – 0				
Umatilla	Minthorn Springs	07_MGILCS	OmyUMA15	2015	53	0 – 159	0	0 – 0				
Lyons Ferry	Mixed origins (Snake + UpC)	07_MGILCS	OmyLYON15S	2015	2,659	1469 – 3842	0	0 – 0	163	0 – 413	0	0 – 0
Little Sheep Creek	-	07_MGILCS	OmyLSCR15S	2015	696	129 – 1341	0	0 – 0				
Wallowa	-	07_MGILCS	OmyWALL15S	2015	4,403	2866 – 6102	58	0 – 173				
Lyons Ferry	Wallowa	07_MGILCS	OmyWALW15S	2015	321	0 – 885	0	0 – 0				
Round Butte	Deschutes River	07_MGILCS	OmyRB16	2016	657	241 – 1193	0	0 – 0				
Lyons Ferry	Mixed origins (Snake + UpC)	07_MGILCS	OmyLYON16S	2016	5,199	3622 – 6804	0	0 – 0	458	36 – 1053	0	0 – 0
Little Sheep Creek	-	07_MGILCS	OmyLSCR16S	2016	301	54 – 615	0	0 – 0				
Wallowa	-	07_MGILCS	OmyWALL16S	2016	647	223 – 1122	0	0 – 0				
Winthrop NFH	Methow River	09_UPPCOL	OmyWTP14	2014	272	0 – 816	0	0 – 0				
Wells	Okanogan Stock	09_UPPCOL	OmyWEL_OKA15	2015	2,383	1291 – 3564	0	0 – 0	133	0 – 398	0	0 – 0
Wells	Omak Stock	09_UPPCOL	OmyWEL_OMA15	2015	610	0 – 1220	0	0 – 0				
Winthrop NFH	Methow River	09_UPPCOL	OmyWTP15	2015	139	0 – 418	0	0 – 0				
Wells	On Station	09_UPPCOL	OmyWEL16S	2016	287	0 – 862	0	0 – 0				
Eastbank	Chelan/Methow/Okanogan/Wenatchee	09_UPPCOL	OmyEASTBK16	2016					133	0 – 398	0	0 – 0
Dworshak NFH	Clearwater River	10_SFCLWR	OmyDWOR14S_1	2014	0	0 – 0	220	0 – 444				
Dworshak NFH	Clearwater River	10_SFCLWR	OmyDWOR14S	2014	0	0 – 0	166	0 – 388				
Dworshak NFH	Clearwater River	10_SFCLWR	OmyDWOR15S	2015	2,636	1873 – 3463	18,508	17028 – 19905	538	238 – 883	3,819	3025 – 4694
Dworshak NFH	Clearwater River	10_SFCLWR	OmyDWOR16S	2016	584	130 – 1173	0	0 – 0	37	0 – 112	0	0 – 0
Oxbow	-	14_UPSALM	OmyOXBO15S	2015	3,319	2045 – 4728	114	0 – 342				
Pahsimeroi	Salmon River	14_UPSALM	OmyPAHH15S	2015	1,786	787 – 2916	196	0 – 482	306	62 – 612	0	0 – 0
Sawtooth	Salmon River	14_UPSALM	OmySAWT15S	2015	257	0 – 572	0	0 – 0				
Oxbow	-	14_UPSALM	OmyOXBO16S	2016	4,274	2730 – 5845	121	0 – 354				
Pahsimeroi	Salmon River	14_UPSALM	OmyPAHH16S	2016	3,341	2185 – 4558	0	0 – 0	192	0 – 504	0	0 – 0
Sawtooth	Salmon River	14_UPSALM	OmySAWT16S	2016	2,784	1665 – 4109	0	0 – 0				
#N/A	#N/A	#N/A	Unassigned	#N/A	4,454	2702 – 6422	265	109 – 582				

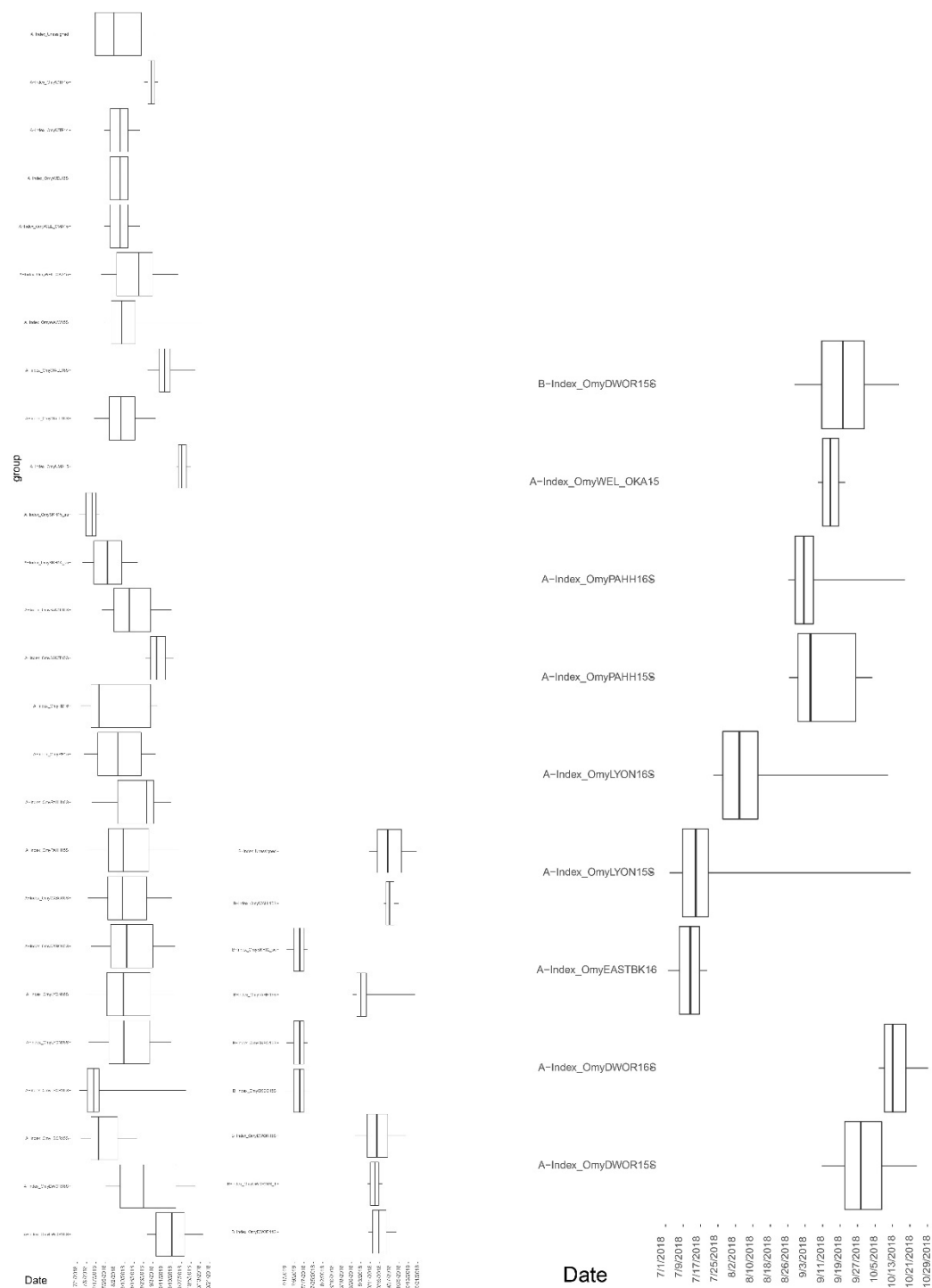
1799 **Note:** These abundance estimates were calculated using a method to estimate abundance of each stock based on temporally stratified stock proportions and total numbers of clipped and unclipped steelhead that
1800 passed the Bonneville Dam at the fish counting window. Key to broodstock collection is presented in Appendix 7.

Run-timing of steelhead stocks in 2018

We were able to characterize the run-timing distributions for hatchery origin clipped and unclipped steelhead stocks (Figure 31). Very few winter-run steelhead stocks exist above Bonneville Dam and our sampling program at Bonneville AFF does not trap or collect fish between December and March when winter-run steelhead would be most likely to occur. The Skamania summer steelhead period is 4/1/2018 – 6/30/ 2018, and the summer A-/B-Index period begins on 7/1/2018 and lasts until 10/31/2018. We examined run-timing of the stocks arriving in the A-/B-Index period (Figure 31). The broodstock that typically has late run-timing is the Dworshak stock, which often arrives after August 25th at Bonneville Dam (Figure 31).

We characterized the run-timing distributions for natural-origin steelhead stocks (Figure 32); the patterns generally are consistent with past years. The late arriving stocks with median dates on or after August 25th were 10_SFCLWR 11_UPCLWR, 13_MFSALM, and 14_UPSALM. We characterized run-timing by A-Index and B-Index categories for these stocks. For some stocks that had earlier run-timing as A-Index sizes, these stocks were later arriving as B-Index stocks (Figure 32).

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Figure 31. Run-timing distributions (median ordinal day, interquartile range, and 5th and 95th percentile) for hatchery-origin steelhead (adipose clipped, left; adipose unclipped, right) that were sampled at Bonneville Dam in 2018 and assigned to broodstock of origin. Each broodstock is shown by A-Index and B-Index size category. Key to broodstock collection is presented in Appendix 7.

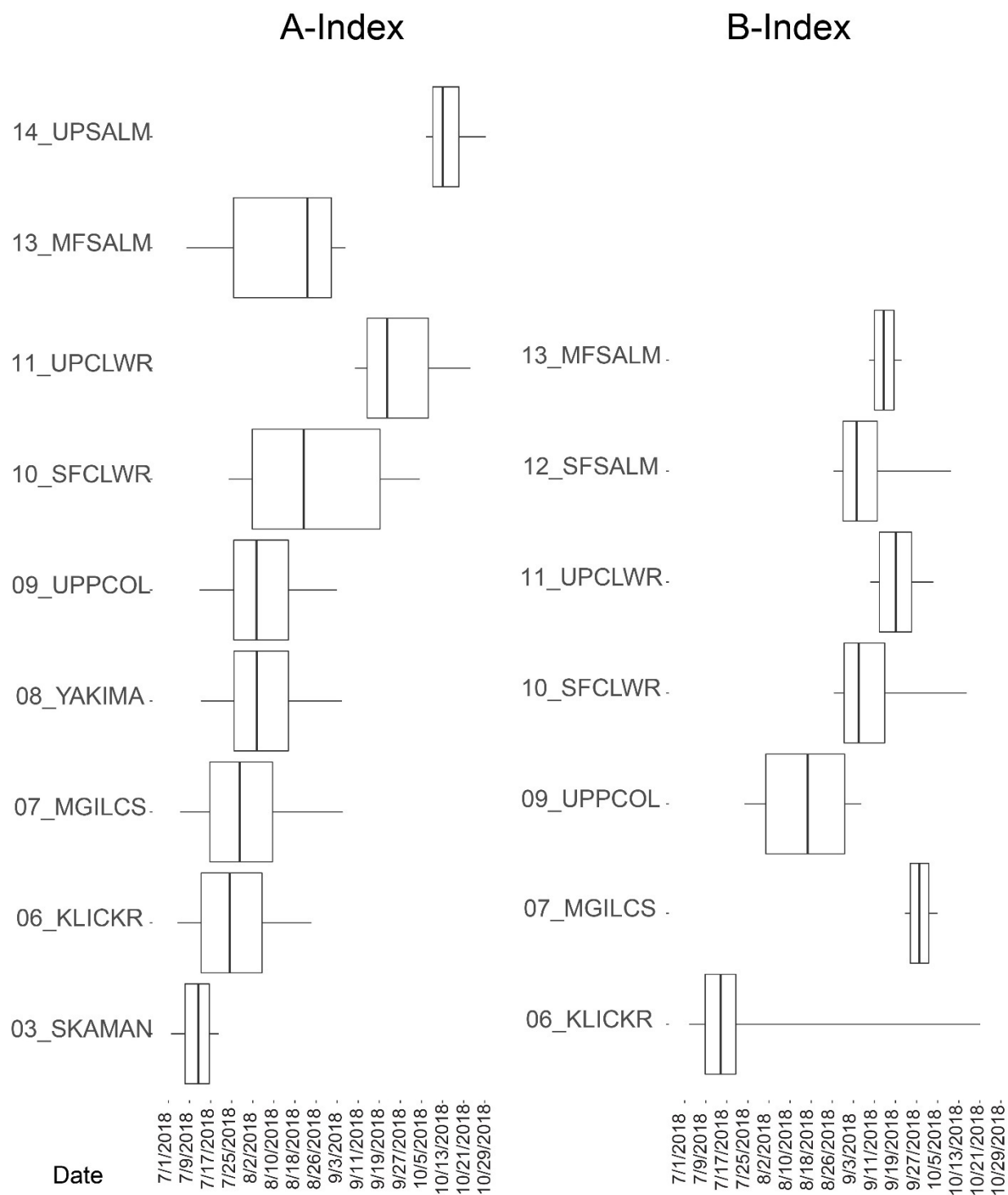


Figure 32. Run-timing distributions (median ordinal day, interquartile range, and 5th and 95th percentile) for natural-origin steelhead (all hatchery-origin adipose-unclipped fish excluded) that were sampled at Bonneville Dam in 2018 and assigned to stock of origin. Each stock is shown by A-Index (left) and B-Index (right) size category

Estimated abundance and run-timing of Sockeye salmon stocks in 2018

Daily passage of Sockeye salmon at Bonneville Dam in 2018 is provided in **Figure 33**. Stock abundance for sockeye salmon was estimated over a course of 16 statistical weeks (i.e. weeks 21-36). A total of 1,857 sockeye salmon were sampled at Bonneville Dam in 2018 and were assigned to one of four genetic stocks (i.e., Okanogan, Wenatchee, Snake, and Lake Billy Chinook) using GSI and one reintroduced stock using PBT (Table 55). The Okanogan stock had the highest relative abundance (174,130), followed by the Wenatchee (17,675) (Table 55). The Snake and Lake Billy Chinook stocks had estimated abundances < 500, but were based on relatively few genetic assignments (<15) (Table 55, **Figure 34**). The reintroduced stock from Yakima River was also low sample size but we estimated 1,294 fish in 2018.

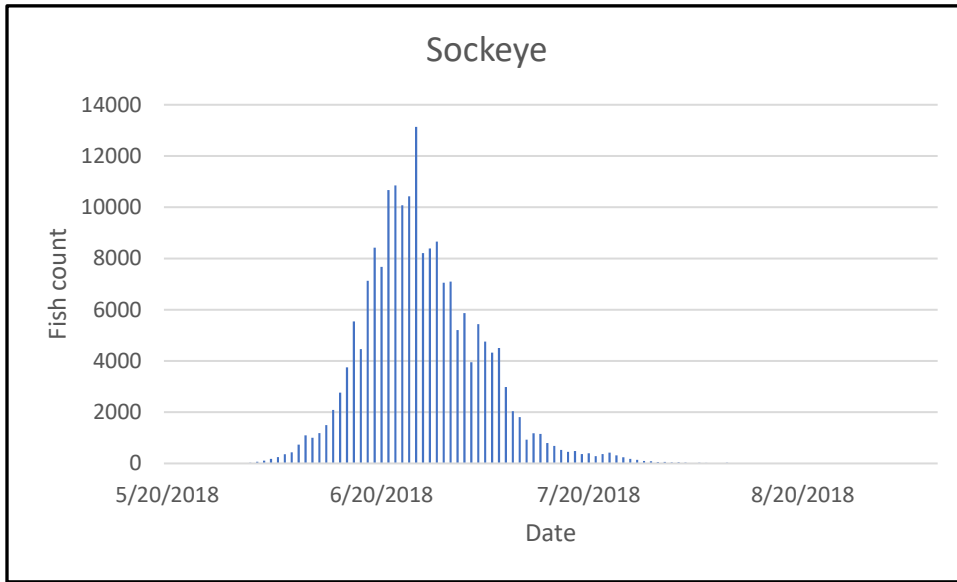


Figure 33. Daily passage of sockeye at Bonneville Dam in 2018 (source: www.fpc.org).

We characterized the run-timing distributions for the two major sockeye salmon stocks (Table 55, Figure 35). The Wenatchee and Okanogan stocks had nearly identical run timing distributions each with a median date near 06/24/18 (Figure 35).

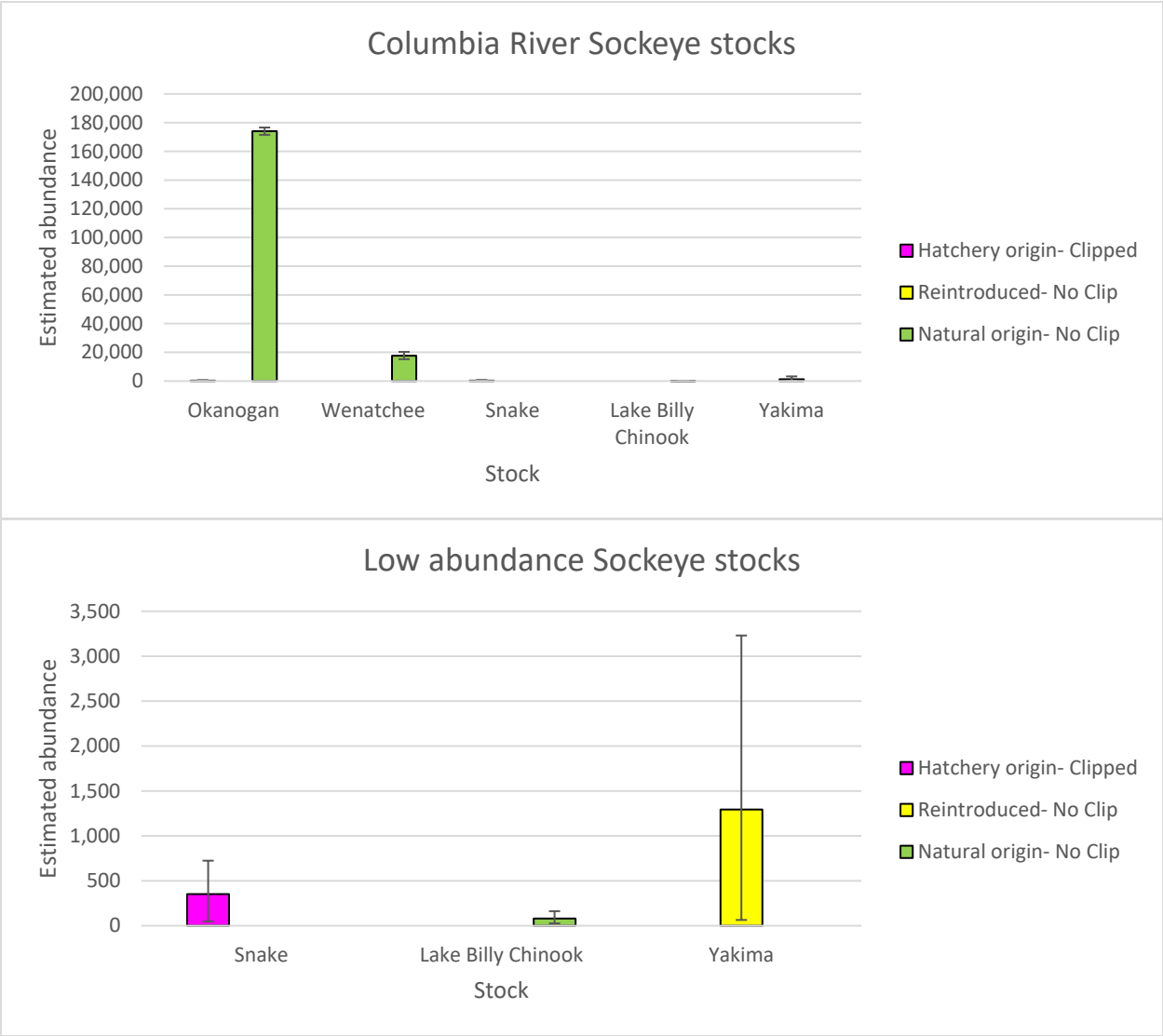


Figure 34. Estimated abundance (\pm 95% CI) of sockeye salmon stocks sampled at Bonneville Dam in 2018.

1850 **Table 55. Estimated abundance of sockeye salmon stocks passing Bonneville Dam in 2018.**

Reporting Group name	Estimated abundance					
	Hatchery origin-Clipped		Reintroduction- No Clip		Natural origin- No Clip	
	Mean	95% CI	Mean	95% CI	Mean	95% CI
Okanogan	285	0 – 630			174,130	171500 – 176606
Wenatchee					17,675	15190 – 20307
Snake	351	49 – 724				
Lake Billy Chinook					80	26 – 162
Yakima			1,294	64 – 3231		

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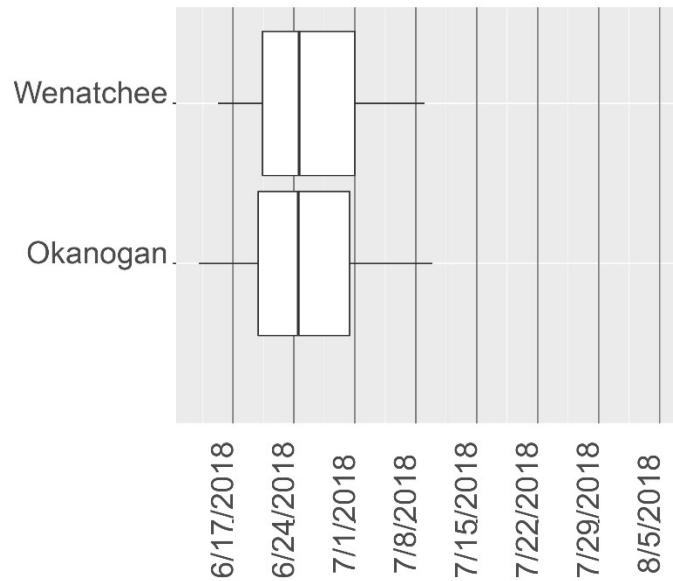


Figure 35. Run-timing distributions (median ordinal day, interquartile range, and 5th and 95th percentile) for the major stocks of sockeye salmon that were sampled at Bonneville Dam in 2018 and assigned to stock of origin.

In-season analysis of Chinook salmon passing Bonneville Dam in 2019

There were four in-season reports covering data on Chinook Salmon that passed Bonneville Dam during the Spring Management Period (Table 47), and a total of seven in-season reports for Chinook Salmon in 2019 across all management periods (Table 56). The first report was distributed to members of the USvOR TAC on May 10, 2019 and included an analysis of data on Chinook Salmon sampled at the Bonneville Dam AFF from the date on which the first Chinook Salmon was sampled (April 25, 2019) until May 3, 2019 (Table 47). We had originally planned to offer reports on approximately a bi-weekly reporting schedule after mid-April when sampling typically begins at the Bonneville Dam AFF, however, the 2019 run of Spring Chinook Salmon was delayed relative to the 10 year average-timed run. The goal for analysis is to have at least two weeks of sample sizes greater than 50 fish per week before in-season reporting is initiated. Each report used a new analysis similar to the reporting for the Bonneville Dam post-season report for run year 2018. However, there was one important change to the post-season methods that we implemented in-season to improve the suitability of the results for managers. Specifically, we used window counts of only the adult-sized fish in order to estimate stock-specific abundances of adult-size Chinook Salmon. In the past, our BPA reports included stock-specific abundance and run-timing estimates of all Chinook Salmon, including jack-sized fish. Management of Chinook Salmon fisheries in the Columbia River is based solely on adult-sized Chinook Salmon (>560 mm fork length), and so we restricted our sample to this fork length threshold for our in-season analysis. Further, we used TAC estimates of the clipped and unclipped adult Chinook salmon and expanded genetic stock proportions with those estimates.

A total of 3,148 adult-sized Chinook Salmon were collected and analyzed for the 2019 in-season reports (Table 56). Estimates of stock-specific abundances of natural-origin stocks (i.e., those fish that were adipose unclipped and did not have a PBT-assignment) and hatchery-origin stocks (i.e. adipose-clipped fish and/or those fish that were PBT-assigned) were provided in each in-season report. Each subsequent report provided cumulative stock-specific abundances and the final report issued on November 1, 2019 provided sub-totals for each stock that were broken out by management period (hatchery-origin stocks, Table 57; natural-origin stocks, Table 59). Two of the stocks that were of particular interest this year were the natural-origin Snake River spring/summer run (Reporting groups 11_TUCANO, 12_HELLSC, 13_SFSALM, 14_CHMBLN, 15_MFSALM, and 16_UPSALM; Table 59) and upper Columbia River spring Chinook Salmon stocks (Reporting group 10_UCOLSP, Table 59). These ESA listed stocks can affect the overall harvest rates that pertain to the Treaty and Non-Treaty fisheries on the mainstem Columbia River. Another aspect of this in-season analysis that was useful to managers was the ability to determine the approximate abundance of stream-type lineage stocks (the spring-run and Snake River spring/summer run stocks) that continue to pass Bonneville Dam during the summer management period. Further, there are non-ESA listed upper Columbia River summer-run (reporting group 18_UCOLSF) that can pass Bonneville Dam before the summer management period. The delineation of these management periods is known by managers to not provide complete separation of these mixed stocks (i.e., interior Columbia River stream-type versus ocean-type genetic lineages), however the dates of the management periods are the result of policy decisions that in part consider a trade-off between over- and under-utilization of ESA and non-ESA listed Chinook Salmon stocks. This in-season genetic analysis can be used in addition to other data by managers to help evaluate options for shaping the fisheries in a way that balances objectives of the USvOR Management Agreement.

PBT assignments during the Chinook Salmon management periods allowed classification of 80 unique hatchery broodstocks in 2019 (Table 60). These broodstocks were comprised of 20% of hatcheries from the Snake River and 75% of hatcheries from the rest of the Columbia River above Bonneville Dam (Figure 36). Similar to the natural- and hatchery-origin abundance estimates, the subtotals of these hatchery broodstock abundances were provided to USvOR TAC for each management period and bi-weekly strata for the in-season reporting in 2019. We observed that this run year of Chinook Salmon at Bonneville Dam was one of the best in terms of coverage of hatchery broodstock in the PBT baseline. One measure of the level of coverage of the PBT baseline is the percentage of clipped adult-sized Chinook Salmon that were estimated to belong to a PBT hatchery broodstock. If all hatchery broodstock above Bonneville Dam are included in the PBT baseline AND our estimated tag rates are accurate for each hatchery broodstock, then this percentage is expected to be 100% across all strata in the all the management periods (Table 61). In fact, in 2019, the percentage of adipose-clipped fish abundance that were estimated to be PBT broodstock ranged from 85-100%, 94-100%, and 82-100% in the spring, summer, and fall management periods, respectively. One reason that the early fall management period had some of the lowest estimated percentages of PBT-assigned clipped fish was likely due to the fact that a major hatchery component of the run during this time, Spring Creek Hatchery, has only recently been added to the PBT baseline and not all broodyears are covered yet (Appendix 6). We expect that even the fall management period will have higher percentages in the future.

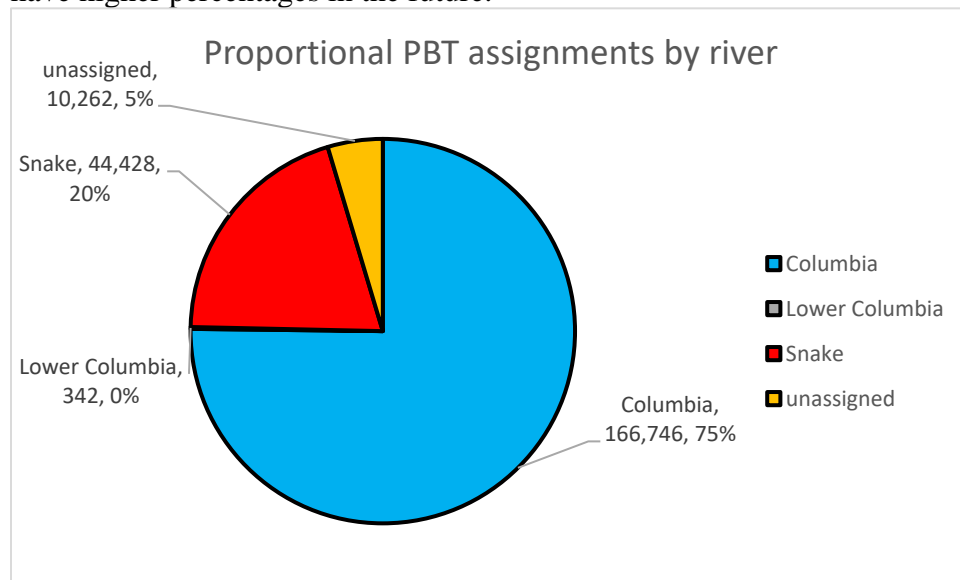


Figure 36. Proportion of hatchery broodstocks from the Snake River versus the rest of the Columbia River above Bonneville Dam in 2019.

1925 **Table 56. The sample sizes of Chinook salmon at the Bonneville Dam AFF during the**
 1926 **spring, summer, and fall management periods of 2019.**

	Statistic al week	TAC- Estimate Clipped Fish	TAC- Estimate Unclipped Fish	Fish count	Sample (N)					Sampl e rate		
					Clipped		Non- clipped		Tota l			
					GS I	PBT	GSI	PB T				
Spring	1-17	1,867	228	2,095	1	10	2	3	16	0.76%		
	18	11,741	2,606	14,347	42	215	38	20	315	2.20%		
	19	14,090	4,357	18,447	19	193	58	13	283	1.53%		
	20	6,268	3,269	9,537	20	95	35	22	172	1.80%		
	21	4,144	2,012	6,156	14	91	43	18	166	2.70%		
	22	3,316	1,621	4,937	8	58	26	9	101	2.05%		
	23	4,922	1,618	6,540	19	63	27	7	116	1.77%		
	24	7,107	2,064	9,171	8	97	27	9	141	1.54%		
Summer	25	7278	1276	8,554	14	67	14	4	99	1.16%		
	26	6502	1590	8,092	8	62	16	4	90	1.11%		
	27	5193	1374	6,567	5	34	22	2	63	0.96%		
	28	3150	972	4,122	4	51	25	4	84	2.04%		
	29	2044	1128	3,172	3	35	22		60	1.89%		
	30	1496	1245	2,741	1	17	13	2	33	1.20%		
	31	501	724	1,225	1	9	17	1	28	2.29%		
Fall	31	412	629	1,041		6	7		13	1.25%		
	32	781	1,663	2,444					0	0.00%		
	33	1,630	3,320	4,950		5	12	4	21	0.42%		
	34	4,405	6,997	11,402	3	24	44	9	80	0.70%		
	35	12,083	18,788	30,871	3	22	54	11	90	0.29%		
	36	17,106	30,167	47,273	6	34	57	14	111	0.23%		
	37	23,583	45,451	69,034	31	81	106	36	254	0.37%		
	38	16,304	27,628	43,932	20	69	178	37	304	0.69%		
	39	10,371	20,080	30,451	12	57	150	28	247	0.81%		
	40	6,403	8,569	14,972	1	10	28	6	45	0.30%		
	41	2,089	5,784	7,873		22	93	20	135	1.71%		
	42-48	3,137	8,686	11,823	2	14	58	7	81	0.69%		
				381,76	24	1,44	1,17		3,14			
Total				177,923	203,846	9	5	1	2	290	8	0.82%

1927 Note: The fish counts indicate the number of adult-sized Chinook Salmon at the fish ladder
 1928 windows at Bonneville Dam and the sample (N) indicates the numbers of adult-sized Chinook
 1929 Salmon (>560 mm fork length) that were collected at the AFF. TAC provides estimates of the
 1930 total clipped an unclipped adult abundance. The AFF sample is broken into adipose-clipped and
 1931 non-clipped categories and then further indicate whether a PBT assignment (PBT) was

1932 confirmed or if it was not assigned with PBT (GSI). Sample rate relates the total sample for a
1933 particular stratum to the total fish counted at the window. The alternating gray and clear rows
1934 indicate the breakpoints in the weekly strata at which time one of the seven total in-season
1935 analyses was performed (Table 47).

1936 **Table 57. Preliminary in-season reporting of clipped hatchery-origin stock-specific abundance estimates of Chinook Salmon**
1937 **passing Bonneville Dam across all management periods in 2019.**

Reporting Group name	Run type	Reporting Group Code	Hatchery origin (Clipped)		Spring		Summer		Fall		Cumulative total date Dec 31
			Estimated abundance		Estimated abundance		Estimated abundance		Estimated abundance		Estimated abundance
			Mean	95% CI	Mean	95% CI	Mean	95% CI	Mean	95% CI	Mean
Youngs Bay	Spring	01_YOUNGS	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0
West Cascade Spring	Spring	02_WCASSP	932	530 – 1368	0	0 – 0	0	0 – 0	0	0 – 0	932
West Cascade Fall*	Fall	03_WCASFA	0	0 – 0	0	0 – 0	763	0 – 2111	763	0 – 2111	763
Willamette	Spring	04_WILLAM	0	0 – 0	0	0 – 251	0	0 – 0	0	0 – 0	0
Spring Creek Tule	Fall	05_SPCRTU	0	0 – 0	0	0 – 0	27,986	21706 – 34298	27,986	21706 – 34298	27,986
Klickitat	Spring	06_KLICKR	73	0 – 218	0	0 – 0	0	0 – 0	0	0 – 0	73
Deschutes spring	Spring	07_DESCSP	577	278 – 900	0	0 – 0	0	0 – 0	0	0 – 0	577
John Day	Spring	08_JOHNDR	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0
Yakima	Spring	09_YAKIMA	941	554 – 1361	0	0 – 0	0	0 – 0	0	0 – 0	941
Upper Columbia spring	Spring	10_UCOLSP	5,926	5061 – 6903	0	0 – 0	0	0 – 0	0	0 – 0	5,926
Tucannon	Spring	11_TUCANO	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0
Hells Canyon	Spring/Summer	12_HELLSC	18,639	17297 – 19844	270	0 – 855	0	0 – 0	0	0 – 0	18,909
South Fork Salmon	Spring/Summer	13_SFSALM	2,442	1801 – 3065	335	0 – 948	0	0 – 0	0	0 – 0	2,777
Chamberlain Creek	Spring/Summer	14_CHMBLN	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0
Middle Fork Salmon	Spring/Summer	15_MFSALM	0	0 – 0	0	0 – 0	0	0 – 0	0	0 – 0	0
Upper Salmon	Spring/Summer	16_UPSALM	2,105	1580 – 2729	92	0 – 454	0	0 – 0	0	0 – 0	2,196

Deschutes fall	Fall	17_DESCFA	0	0 – 0	0	0 – 0	0	0 – 0	0
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	12,976	12317 – 13587	25,167	24163 – 25916	21,153	16605 – 27655	59,296
Snake River fall	Fall	19_SRFALL	0	0 – 0	300	0 – 795	9,958	6182 – 14213	10,258
Bonneville Pool spring	Spring	20_BONPOOLSP	8,493	7429 – 9556	0	0 – 0	0	0 – 0	8,493
Umatilla spring	Spring	21_UMATILLASP	351	136 – 613	0	0 – 0	0	0 – 0	351
Bonneville Pool fall	Fall	22_BONPOOLFA	0	0 – 0	0	0 – 0	38,446	32517 – 45044	38,446
Umatilla fall	Fall	23_UMATILLAFA	0	0 – 0	0	0 – 0	0	0 – 0	0
Total			53,455		26,164		98,305		177,923

Note: the sub-total estimates for each stock are provided for the spring (jan 1 – jun 15), summer (jun 16 – jul 31), and fall (aug 1 – dec 31) management periods and the cumulative total through the window counts on Dec 31, 2019 are provided.

Table 58. Preliminary in-season reporting of unclipped hatchery-origin stock-specific abundance estimates of Chinook Salmon passing Bonneville Dam across all management periods in 2019.

		Hatchery origin (No Clip)	Spring		Summer		Fall		Cumulative total date Dec 31
Reporting Group name	Run type	Reporting Group Code	Estimated abundance		Estimated abundance		Estimated abundance		Estimated abundance
			Mean	95% CI	Mean	95% CI	Mean	95% CI	Mean
Youngs Bay	Spring	01_YOUNGS	0	0 – 0	0	0 – 0	0	0 – 0	0
West Cascade Spring	Spring	02_WCASSP	34	0 – 103	0	0 – 0	0	0 – 0	34
West Cascade Fall*	Fall	03_WCASFA	0	0 – 0	0	0 – 0	0	0 – 0	0
Willamette	Spring	04_WILLAM	0	0 – 0	95	0 – 286	0	0 – 0	95
Spring Creek Tule	Fall	05_SPCRTU	0	0 – 0	0	0 – 0	3,869	1079 – 7481	3,869
Klickitat	Spring	06_KLICKR	0	0 – 0	0	0 – 0	0	0 – 0	0

Deschutes spring	Spring	07_DESCSP	117	0 – 286	0	0 – 0	0	0 – 0	117
John Day	Spring	08_JOHNDR	0	0 – 0	0	0 – 0	0	0 – 0	0
Yakima	Spring	09_YAKIMA	34	0 – 102	0	0 – 0	0	0 – 0	34
Upper Columbia spring	Spring	10_UCOLSP	2,634	2005 – 3285	0	0 – 0	0	0 – 0	2,634
Tucannon	Spring	11_TUCANO	127	0 – 292	0	0 – 0	0	0 – 0	127
Hells Canyon	Spring/Summer	12_HELLSC	885	541 – 1245	80	0 – 240	0	0 – 0	965
South Fork Salmon	Spring/Summer	13_SFSALM	793	462 – 1150	0	0 – 0	0	0 – 0	793
Chamberlain Creek	Spring/Summer	14_CHMBLN	0	0 – 0	0	0 – 0	0	0 – 0	0
Middle Fork Salmon	Spring/Summer	15_MFSALM	0	0 – 0	0	0 – 0	0	0 – 0	0
Upper Salmon	Spring/Summer	16_UPSALM	0	0 – 0	0	0 – 0	0	0 – 0	0
Deschutes fall	Fall	17_DESCFA	0	0 – 0	0	0 – 0	0	0 – 0	0
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	497	256 – 746	863	532 – 1230	14,900	10588 – 19209	16,260
Snake River fall	Fall	19_SRFALL	0	0 – 0	64	0 – 192	10,315	6247 – 15066	10,379
Bonneville Pool spring	Spring	20_BONPOOLSP	191	46 – 342	0	0 – 0	0	0 – 0	191
Umatilla spring	Spring	21_UMATILLASP	632	343 – 954	0	0 – 0	0	0 – 0	632
Bonneville Pool fall	Fall	22_BONPOOLFA	0	0 – 0	0	0 – 0	7,697	4554 – 10950	7,697
Umatilla fall	Fall	23_UMATILLAFA	0	0 – 0	0	0 – 0	0	0 – 0	0
Total			5,944		1,103		36,781		43,827

1945 Note: the sub-total estimates for each stock are provided for the spring (jan 1 – jun 15), summer (jun 16 – jul 31), and fall (aug 1 – dec
1946 31) management periods and the cumulative total through the window counts on Dec 31, 2019 are provided.
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1949 **Table 59. Preliminary in-season reporting of natural-origin stock-specific abundance estimates of Chinook Salmon passing**
1950 **Bonneville Dam across all management periods in 2019**

		Natural origin- No Clip	Spring		Summer		Fall		Cumulative total date Dec 31
Reporting Group name	Run type	Reporting Group Code	Estimated abundance		Estimated abundance		Estimated abundance		Estimated abundance
			Mea n	95% CI	Mea n	95% CI	Mea n	95% CI	Mean
Youngs Bay	Spring	01_YOUNGS	0	0 – 0	0	0 – 0	0	0 – 0	0
West Cascade Spring	Spring	02_WCASSP	0	0 – 0	0	0 – 0	0	0 – 0	0
West Cascade Fall*	Fall	03_WCASFA	0	0 – 0	0	0 – 0	1,271	478 – 2256	1,271
Willamette	Spring	04_WILLAM	0	0 – 0	0	0 – 0	0	0 – 0	0
Spring Creek Tule	Fall	05_SPCRTU	0	0 – 0	0	0 – 0	1,214	408 – 2934	1,214
Klickitat	Spring	06_KLICKR	0	0 – 0	0	0 – 0	0	0 – 0	0
Deschutes spring	Spring	07_DESCSP	468	226 – 748	0	0 – 0	0	0 – 0	468
John Day	Spring	08_JOHNDR	674	401 – 976	0	0 – 0	0	0 – 0	674
Yakima	Spring	09_YAKIMA	928	609 – 1295	51	0 – 154	0	0 – 0	979
Upper Columbia spring	Spring	10_UCOLSP	1,712	1207 – 2328	0	0 – 0	0	0 – 0	1,712
Tucannon	Spring	11_TUCANO	91	0 – 183	0	0 – 0	0	0 – 0	91
Hells Canyon	Spring/Su mmer	12_HELLSC	3,584	2941 – 4225	78	0 – 235	0	0 – 0	3,662
South Fork Salmon	Spring/Su mmer	13_SFSALM	779	455 – 1109	78	0 – 236	0	0 – 0	857
Chamberlain Creek	Spring/Su mmer	14_CHMBLN	0	0 – 0	0	0 – 0	0	0 – 0	0
Middle Fork Salmon	Spring/Su mmer	15_MFSALM	353	151 – 591	0	0 – 0	0	0 – 0	353
Upper Salmon	Spring/Su mmer	16_UPSALM	914	604 – 1252	57	0 – 172	0	0 – 0	971

Deschutes fall	Fall	17_DESCFA	0	0 – 0	0	0 – 0	2,801	1706 – 4128	2,801
Upper Columbia summer/fall	Summer/Fall	18_UCOLSF	2,273	1824 – 2680	6,653	6158 – 7137	115,574	109979 – 120502	124,500
Snake River fall	Fall	19_SRFALL	55	0 – 164	289	93 – 527	20,120	16799 – 23652	20,464
Total			11,831		7,207		140,980		160,018

Note: the sub-total estimates for each stock are provided for the spring (jan 1 – jun 15), summer (jun 16 – jul 31), and fall (aug 1 – dec 31) management periods and the cumulative total through the window counts on Dec 31, 2019 are provided.

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1958 **Table 60. The estimated abundances of the clipped and unclipped PBT hatchery broodstock that passed Bonneville Dam in**
1959 **2019 (1/1/2019 – 12/31/2019)**

Expected Run Time	Hatchery	Broodstock	Brood year	Lineage	GSI RepGrp	Snake / Columbia	MLE		
							Clipped	Unclipped	Total
01Spring	Parkdale Fish Facility	OtsPFF14	2014	Lower Columbia	02_WCASSP	Columbia	70	0	70
01Spring	Parkdale Fish Facility	OtsPFF15	2015	Lower Columbia	02_WCASSP	Columbia	772	34	807
01Spring	South Santiam Hatchery	OtsSSANT1 5S	2015	Lower Columbia	04_WILLAM	Lower Columbia	0	86	86
01Spring	Klickitat State Fish Hatchery	OtsKH14	2014	Interior stream type	06_KLICKR	Columbia	71	0	71
01Spring	Round Butte Fish Hatchery	OtsRB15	2015	Interior stream type	07_DESCSP	Columbia	424	117	541
01Spring	Warm Springs National Fish Hatchery	OtsWSNFH 15	2015	Interior stream type	07_DESCSP	Columbia	501	0	501
01Spring	Levi George/Cle Elum (Integrated)	OtsYR15int	2015	Interior stream type	09_YAKIMA	Columbia	722	34	756
01Spring	Levi George/Cle Elum (Segregated)	OtsYR15seg	2015	Interior stream type	09_YAKIMA	Columbia	125	0	125
01Spring	Chief Joseph Hatchery (Spring)	OtsCJH15_s p	2015	Interior stream type	10_UCOLSP	Columbia	1,666	58	1,723
01Spring	Eastbank Fish Hatchery (Spring)	OtsEASTBK 14_sp	2014	Interior stream type	10_UCOLSP	Columbia	0	86	86
01Spring	Eastbank Fish Hatchery (Spring)	OtsEASTBK 15_sp	2015	Interior stream type	10_UCOLSP	Columbia	468	1,047	1,515
01Spring	Eastbank Fish Hatchery (Spring)	OtsEASTBK 16S_sp	2016	Interior stream type	10_UCOLSP	Columbia	0	98	98
01Spring	Leavenworth National Fish Hatchery	OtsLNFH15	2015	Interior stream type	10_UCOLSP	Columbia	1,772	53	1,825
01Spring	Methow State Fish Hatchery	OtsMETH14	2014	Interior stream type	10_UCOLSP	Columbia	0	46	46
01Spring	Methow State Fish Hatchery	OtsMETH15	2015	Interior stream type	10_UCOLSP	Columbia	0	411	411

01Spring	Winthrop National Fish Hatchery	OtsWTP15	2015	Interior stream type	10_UCOLSP	Columbia	1,764	835	2,599
01Spring	Lyons Ferry Fish Hatchery (Spring)	OtsLYON15S	2015	Interior stream type	11_TUCANO	Snake	0	127	127
01Spring	Clearwater Fish Hatchery	OtsCLWH14S	2014	Interior stream type	12_HELLSC	Snake	293	0	293
01Spring	Clearwater Fish Hatchery	OtsCLWH15S	2015	Interior stream type	12_HELLSC	Snake	2,891	198	3,089
01Spring	Clearwater Fish Hatchery - Powell Facility	OtsPOWP14S	2014	Interior stream type	12_HELLSC	Snake	114	0	114
01Spring	Clearwater Fish Hatchery - Powell Facility	OtsPOWP16S	2016	Interior stream type	12_HELLSC	Snake	0	86	86
01Spring	Dworshak National Fish Hatchery	OtsDWOR14S	2014	Interior stream type	12_HELLSC	Snake	86	0	86
01Spring	Dworshak National Fish Hatchery	OtsDWOR15S	2015	Interior stream type	12_HELLSC	Snake	2,808	164	2,972
01Spring	Lookingglass Fish Hatchery - Catherine Creek	OtsCTHW15S	2015	Interior stream type	12_HELLSC	Snake	449	0	449
01Spring	Lookingglass Fish Hatchery - Grande Ronde	OtsGRUW15S	2015	Interior stream type	12_HELLSC	Snake	257	231	488
01Spring	Lookingglass Fish Hatchery - Lookingglass Creek	OtsLOOK14S	2014	Interior stream type	12_HELLSC	Snake	49	0	49
01Spring	Lookingglass Fish Hatchery - Lookingglass Creek	OtsLOOK15S	2015	Interior stream type	12_HELLSC	Snake	676	0	676
01Spring	Lookingglass Fish Hatchery - Lostine River	OtsLSTW14S	2014	Interior stream type	12_HELLSC	Snake	54	0	54
01Spring	Lookingglass Fish Hatchery - Lostine River	OtsLSTW15S	2015	Interior stream type	12_HELLSC	Snake	250	0	250
01Spring	Rapid River Fish Hatchery	OtsRAPH14S	2014	Interior stream type	12_HELLSC	Snake	377	0	377
01Spring	Rapid River Fish Hatchery	OtsRAPH15S	2015	Interior stream type	12_HELLSC	Snake	8,592	253	8,845

01Spring	Carson National Fish Hatchery	OtsCAR14	2014	Interior stream type	20_BONPOOL SP	Columbia	239	0	239
01Spring	Carson National Fish Hatchery	OtsCAR15	2015	Interior stream type	20_BONPOOL SP	Columbia	3,396	139	3,535
01Spring	Little White Salmon National Fish Hatchery (Spring)	OtsLWS14_sp	2014	Interior stream type	20_BONPOOL SP	Columbia	605	0	605
01Spring	Little White Salmon National Fish Hatchery (Spring)	OtsLWS15_sp	2015	Interior stream type	20_BONPOOL SP	Columbia	3,865	52	3,917
01Spring	Umatilla Fish Hatchery (Spring)	OtsUMA14_sp	2014	Interior stream type	21_UMATILL ASP	Columbia	67	0	67
01Spring	Umatilla Fish Hatchery (Spring)	OtsUMA15_sp	2015	Interior stream type	21_UMATILL ASP	Columbia	244	632	876
02Spring/Summer	Lookingglass Fish Hatchery - Imnaha River	OtsIMNW14S	2014	Interior stream type	12_HELLSC	Snake	43	0	43
02Spring/Summer	Lookingglass Fish Hatchery - Imnaha River	OtsIMNW15S	2015	Interior stream type	12_HELLSC	Snake	430	38	468
02Spring/Summer	Lookingglass Fish Hatchery - Imnaha River	OtsIMNW16S	2016	Interior stream type	12_HELLSC	Snake	194	0	194
02Spring/Summer	McCall Fish Hatchery - South Fork Salmon	OtsMCCA15S	2015	Interior stream type	13_SFSALM	Snake	2,510	793	3,302
02Spring/Summer	McCall Fish Hatchery - South Fork Salmon	OtsMCCA16S	2016	Interior stream type	13_SFSALM	Snake	228	0	228
02Spring/Summer	Pahsimeroi Fish Hatchery	OtsPAHH14S	2014	Interior stream type	16_UPSALM	Snake	46	0	46
02Spring/Summer	Pahsimeroi Fish Hatchery	OtsPAHH15S	2015	Interior stream type	16_UPSALM	Snake	612	0	612
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT14S	2014	Interior stream type	16_UPSALM	Snake	176	0	176
02Spring/Summer	Sawtooth Fish Hatchery	OtsSAWT15S	2015	Interior stream type	16_UPSALM	Snake	1,068	0	1,068
03Summer	Chief Joseph Hatchery	OtsCJH14int_su	2014	Interior ocean type	18_UCOLSF	Columbia	513		513
03Summer	Chief Joseph Hatchery	OtsCJH15int_su	2015	Interior ocean type	18_UCOLSF	Columbia	813		813

03Summer	Chief Joseph Hatchery	OtsCJH16int _su	2016	Interior ocean type	18_UCOLSF	Columbia	257		257
03Summer	Chief Joseph Hatchery (Summer/Fall) - Integrated	OtsCJH14int _sufa	2014	Interior ocean type	18_UCOLSF	Columbia	4,716	224	4,940
03Summer	Chief Joseph Hatchery (Summer/Fall) - Integrated	OtsCJH15int _sufa	2015	Interior ocean type	18_UCOLSF	Columbia	2,388	164	2,552
03Summer	Chief Joseph Hatchery (Summer/Fall) - Integrated	OtsCJH16int _sufa	2016	Interior ocean type	18_UCOLSF	Columbia	346	86	432
03Summer	Chief Joseph Hatchery (Summer/Fall) - Segregated	OtsCJH13se g_sufa	2013	Interior ocean type	18_UCOLSF	Columbia	169	0	169
03Summer	Chief Joseph Hatchery (Summer/Fall) - Segregated	OtsCJH14se g_sufa	2014	Interior ocean type	18_UCOLSF	Columbia	4,211	0	4,211
03Summer	Chief Joseph Hatchery (Summer/Fall) - Segregated	OtsCJH15se g_sufa	2015	Interior ocean type	18_UCOLSF	Columbia	2,021	209	2,230
03Summer	Chief Joseph Hatchery (Summer/Fall) - Segregated	OtsCJH16se g_sufa	2016	Interior ocean type	18_UCOLSF	Columbia	54	0	54
03Summer	Eastbank Fish Hatchery	OtsEASTBK 15_su	2015	Interior ocean type	18_UCOLSF	Columbia	997		997
03Summer	Eastbank Fish Hatchery (Summer)	OtsEASTBK 13_su	2013	Interior ocean type	18_UCOLSF	Columbia	133	0	133
03Summer	Eastbank Fish Hatchery (Summer)	OtsEASTBK 14_su	2014	Interior ocean type	18_UCOLSF	Columbia	6,077	61	6,138
03Summer	Eastbank Fish Hatchery (Summer)	OtsEASTBK 15_su	2015	Interior ocean type	18_UCOLSF	Columbia	4,570	184	4,754
03Summer	Entiat National Fish Hatchery (Summer)	OtsENFH14	2014	Interior ocean type	18_UCOLSF	Columbia	2,076	0	2,076
03Summer	Entiat National Fish Hatchery (Summer)	OtsENFH15	2015	Interior ocean type	18_UCOLSF	Columbia	3,578	0	3,578
03Summer	Wells Fish Hatchery	OtsWELLS1 4	2014	Interior ocean type	18_UCOLSF	Columbia	4,530	109	4,640
03Summer	Wells Fish Hatchery	OtsWELLS1 5	2015	Interior ocean type	18_UCOLSF	Columbia	2,470	343	2,813

04Fall	Washougal Fish Hatchery	OtsWAS15_fa	2015	Lower Columbia	03_WCASFA	Lower Columbia	257		257
04Fall	Spring Creek National Fish Hatchery	OtsSPCR15_fa	2015	Lower Columbia	05_SPCRTU	Columbia	1,083	285	1,368
04Fall	Spring Creek National Fish Hatchery	OtsSPCR16_fa	2016	Lower Columbia	05_SPCRTU	Columbia	11,235	2,010	13,244
04Fall	Spring Creek National Fish Hatchery	OtsSPCR17_fa	2017	Lower Columbia	05_SPCRTU	Columbia	11,949	1,575	13,524
04Fall	Little White Salmon National Fish Hatchery	OtsLWS14_fa	2014	Interior ocean type	18_UCOLSF	Columbia	835	596	1,431
04Fall	Little White Salmon National Fish Hatchery	OtsLWS15_fa	2015	Interior ocean type	18_UCOLSF	Columbia	23,428	1,911	25,339
04Fall	Little White Salmon National Fish Hatchery	OtsLWS16_fa	2016	Interior ocean type	18_UCOLSF	Columbia	14,183	5,189	19,372
04Fall	Priest Rapids Hatchery	OtsPRH14_fa	2014	Interior ocean type	18_UCOLSF	Columbia		51	51
04Fall	Priest Rapids Hatchery	OtsPRH15_fa	2015	Interior ocean type	18_UCOLSF	Columbia	7,189	6,932	14,121
04Fall	Priest Rapids Hatchery	OtsPRH16_fa	2016	Interior ocean type	18_UCOLSF	Columbia	8,664	7,917	16,581
04Fall	Lyons Ferry Fish Hatchery	OtsLYON14_S_1_fa	2014	Interior ocean type	19_SRFALL	Snake	257	1,033	1,290
04Fall	Lyons Ferry Fish Hatchery	OtsLYON15_S_1_fa	2015	Interior ocean type	19_SRFALL	Snake	4,147	3,012	7,158
04Fall	Lyons Ferry Fish Hatchery	OtsLYON16_S_1_fa	2016	Interior ocean type	19_SRFALL	Snake	3,944	2,612	6,556
04Fall	Lyons Ferry Fish Hatchery (Fall)	OtsLYON15_S_1	2015	Interior ocean type	19_SRFALL	Snake	227	74	301
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH15_S_1_fa	2015	Interior ocean type	19_SRFALL	Snake	613	2,425	3,038
04Fall	Nez Perce Tribal Fish Hatchery	OtsNPFH16_S_1_fa	2016	Interior ocean type	19_SRFALL	Snake	685	1,234	1,919
04Fall	Nez Perce Tribal Fish Hatchery (Fall)	OtsNPFH16_S_1	2016	Interior ocean type	19_SRFALL	Snake	73	0	73
#N/A	#N/A	Unassigned	#N/A	#N/A	#N/A	#N/A	10,262	0	10,262
Grand Total							177,923	43,854	221,778

1961
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Table 61. Expanded abundance of PBT-assigned Chinook Salmon stocks across management periods in 2019

		Sample (Tag-rate-Corrected abundance)						% PBT of Clipped fish	
	Statistical	Clipped		Non-clipped		Total GSI	Total PBT		
	week	GSI	PBT	GSI	PBT				
Management period	Spring	1-17	33	1,833	64	164	98	1,997	98.2%
		18	1,106	10,635	1,591	1,015	2,697	11,650	90.6%
		19	415	13,676	3,492	864	3,907	14,540	97.1%
		20	693	5,575	1,843	1,426	2,535	7,002	89.0%
		21	236	3,908	1,277	736	1,512	4,644	94.3%
		22	-62	3,378	1,057	563	995	3,942	101.9%
		23	728	4,193	1,245	374	1,973	4,567	85.2%
		24	-262	7,369	1,449	615	1,187	7,984	103.7%
	Summer	25	430	6,848	977	299	1,407	7,147	94.1%
		26	-148	6,650	1,247	343	1,099	6,993	102.3%
		27	-166	5,359	1,253	121	1,087	5,480	103.2%
		28	-291	3,441	827	145	536	3,586	109.2%
		29	-167	2,211	1,128	0	961	2,211	108.2%
		30	-77	1,573	1,066	179	989	1,752	105.2%
		31	18	483	681	43	699	526	96.4%
	Fall	31	-28	509	561	0	532	509	105.9%
		32							
		33	-247	635	1,647	882	1,400	1,518	163.7%
		34	-357	1,843	3,463	1,173	3,106	3,016	124.0%
		35	-638	3,994	11,110	2,459	10,473	6,453	119.0%
		36	-134	6,298	14,397	4,899	14,263	11,197	102.2%
		37	1,862	8,537	18,477	6,933	20,339	15,469	82.1%
		38	622	4,151	15,937	3,602	16,560	7,753	87.0%
		39	243	2,654	12,095	2,375	12,339	5,029	91.6%
		40	128	1,437	5,332	1,143	5,460	2,579	91.8%
		41	-22	363	3,934	907	3,912	1,270	106.6%
		42-48	76	544	6,215	756	6,290	1,300	87.8%

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Note: The % PBT of Clipped fish indicates the percentage of adipose clipped fish that are accounted for by PBT after tag rate expansions. If all hatchery broodstock above Bonneville Dam are included in the PBT baseline AND our estimated tag rates are accurate for each hatchery broodstock, then this percentage is expected to be 100% across all strata.
*The % values above 100% can occur when PBT expansion is greater than the total count of clipped fish in a stratum.

In-season analysis of steelhead passing Bonneville Dam in 2019

There were three reports provided to USvOR TAC during the steelhead management periods: Skamania Management Period (4/1/2019 – 6/30/2019) and summer A-/B-Index Management Period (7/1/2019 – 10/31/2019, Table 47). There were a total of 471 adipose clipped and 371 adipose unclipped steelhead that were sampled at the Bonneville Dam AFF and genotyped in 2019 (Table 62). The methods described for the analysis of the 2017 sample of steelhead at Bonneville Dam are not as useful to fisheries managers that serve on USvOR TAC because the size groups (A-/B-Index) have not been fully integrated into the results. Therefore, we revised these methods and provided a breakdown of those A-/B-Index groups in addition to the adipose clipped and unclipped categories (Table 62). One of the important features of the genetic analyses of steelhead at Bonneville Dam is that they can identify unmarked hatchery fish via PBT assignments. The identification of unmarked hatchery fish is critical for accurate calculations of ESA impacts on steelhead stocks. The methods we are using provide a means to estimate the abundance of six categories of steelhead; i.e., clipped hatchery (H), unclipped hatchery (HNC), and natural-origin (W) for A-Index and B-Index fish (Table 63). We also incorporate tag rate expansion to account for the percentage of each hatchery stock that has been successfully genotyped. Tag rate expansions have been used previously in all of our genetic analyses for Chinook Salmon and steelhead. However, one feature that has been missing from our tag rate expansion methods in the past is a way to balance the numbers of fish that are expanded in the PBT broodstock categories by subtracting an equal set of fish from the sample. This issue is now resolved through our implementation of SCOBIDEUX and SPIBETR methodologies described previously (Section 3).

1993
1994

Table 62. The sample sizes of Summer Steelhead at the Bonneville Dam AFF during the Skamania and A-/B-Index management periods of 2019.

		Sample (N)														Clipp ed Samp le rate	Non- clipped Sample rate
		Statistical week	Stra ta	Clipp ed count	Non- Clipped count	A-Index				B-Index				Clipp ed Total	Non- clipped Total		
						Clipped GS I	PB T	Non- clipped GS I	PB T	Clipped GS I	PB T	Non- clipped GS I	PB T				
Management period Skamania	14	1	67	105									0	0	0.00 %	0.00%	
	15	1	53	76									0	0	0.00 %	0.00%	
	16	1	44	46									0	0	0.00 %	0.00%	
	17	1	42	26			1	1					1	1	2.38 %	3.85%	
	18	1	92	28									0	0	0.00 %	0.00%	
	19	1	81	37				1					0	1	0.00 %	2.70%	
	20	1	62	28				2					0	2	0.00 %	7.14%	
	21	1	57	23			1	1					1	1	1.75 %	4.35%	
	22	1	70	56	1	1							2	0	2.86 %	0.00%	
	23	1	129	53			4						4	0	3.10 %	0.00%	
	24	1	106	135				1			1		0	2	0.00 %	1.48%	
	25	1	235	288			1	4					1	4	0.43 %	1.39%	
	26	1	457	738			5	3					5	3	1.09 %	0.41%	

	Subtotal Skamania	1	1495	1639	1	13	13	0	0	0	1	0	14	14	0.94 %	0.85%
A-/B-Index	27	1	1160	1,745	1	5	8				1		6	9	0.52 %	0.52%
	28	1	1445	2,516	4	8	18						12	18	0.83 %	0.72%
	29	1	2205	3,503	4	19	36				2		23	38	1.04 %	1.08%
	30	2	2754	4,089	2	15	25						17	25	0.62 %	0.61%
	31	2	3170	4,167	3	29	37						32	37	1.01 %	0.89%
	32	2	3837	4,529									0	0	0.00 %	0.00%
	33	3	4185	4,051	2	21	33	1					23	34	0.55 %	0.84%
	34	3	2878	1,911	7	33	34				1		40	35	1.39 %	1.83%
	35	4	2779	1,704	5	27	21	1		1			33	22	1.19 %	1.29%
	36	4	2011	1,166	1	12	7	2		1		1	14	10	0.70 %	0.86%
	37	4	2108	1,131	3	16	5	2		2	1	1	21	9	1.00 %	0.80%
	38	5	2960	1,328	2	28	10	3		17	3	3	47	19	1.59 %	1.43%
	39	5	1569	633	1	10	6	2		10	2		21	10	1.34 %	1.58%
	40	6	1366	606	2	19	7	5	1	21	3	11	43	26	3.15 %	4.29%
	41	6	1708	825		36	10	9		30	3	5	66	27	3.86 %	3.27%
	42	7	929	474	1	32	11	7		12	2	4	45	24	4.84 %	5.06%

43	7	510	285											0.00	
														%	0.00%
44	7	147	82											0.00	
														%	0.00%
<hr/>															
Summer A-/B- Index subtotal		37,72		31	26									1.17	
		1	34,745	38	0	8	32	1	94	18	25	443	343	%	0.99%
		40,71		33	29									1.16	
Total		1	38,023	40	6	4	32	1	94	20	25	471	371	%	0.98%

1995 Note: The clipped and non-clipped counts indicate the number of adipose clipped and unclipped steelhead at the fish ladder windows
1996 at Bonneville Dam and the sample (N) indicates the numbers of A- and B-sized (<580 mm and ≥580 mm fork length, respectively)
1997 that were collected at the AFF. The AFF sample is broken into A- and B-sized and adipose-clipped and non-clipped categories and
1998 then further indicate whether a PBT assignment (PBT) was confirmed or if it was not assigned with PBT (GSI). Sample rate relates
1999 the total sample for a particular stratum to the total fish counted at the window. The alternating gray and clear rows indicate the
2000 breakpoints in the weekly strata at which time one of the three total in-season analyses was performed (Table 47).
2001
2002
2003
2004
2005

Table 63. Estimated abundance of hatchery-origin clipped stocks of Summer Steelhead that passed Bonneville Dam in 2019 during the Skamania and A-/B-Index Management Periods.

		Hatchery origin- Clipped	Subtotal for Skamania Period (Apr 1 - Jun 30)		Subtotal for A-/B-Index Period (Jul 01 - Oct 31)		Subtotal for A-/B-Index Period (Jul 01 - Oct 31)	
Reporting Group name	Reporting Group Code	Estimated abundance		A-INDEX: Estimated abundance		B-INDEX: Estimated abundance		
		Mean	95% CI	Mean	95% CI	Mean	95% CI	
Lower Columbia	02_LOWCO L	0	0 – 0	0	0 – 0	0	0 – 0	
Skamania	03_SKAMA N	1,281	854 – 1495	3,082	1480 – 4553	0	0 – 0	
Willamette	04_WILLA M	0	0 – 0	0	0 – 0	0	0 – 0	

Big White Salmon	05_BWSAL M	0	0 – 0	0	0 – 0	0	0 – 0
Klickitat	06_KLICKR	0	0 – 0	0	0 – 0	0	0 – 0
mid Columbia/Grande Ronde/Imnaha/lower Clearwater/lower Snake/lower Salmon	07_MGILCS	0	0 – 0	14,240	11754 – 17247	0	0 – 0
Yakima	08_YAKIM A	0	0 – 0	0	0 – 0	0	0 – 0
upper Columbia	09_UPPCOL	214	0 – 641	990	219 – 1929	0	0 – 0
SF Clearwater	10_SFCLWR	0	0 – 0	3,783	2655 – 4894	4,029	3096 – 4977
upper Clearwater	11_UPCLW R	0	0 – 0	34	0 – 223	0	0 – 0
SF Salmon	12_SFSALM	0	0 – 0	34	0 – 219	0	0 – 0
MF Salmon	13_MFSAL M	0	0 – 0	0	0 – 0	0	0 – 0
upper Salmon	14_UPSALM	0	0 – 0	11,465	8909 – 14418	64	0 – 263
	Total	1,495		33,628		4,093	

Note: Based on the sample data described in Table 62 we estimated abundances of six categories of steelhead; i.e., clipped hatchery (H), unclipped hatchery (HNC), and natural-origin (W) for A-Index and B-Index fish. The sample was divided into the most strata possible maintaining a minimum sample of 100 clipped and unclipped steelhead per stratum in the A-/B-Index period. All weeks were pooled into a single stratum in the Skamania period.

Table 64. Estimated abundance of hatchery-origin unclipped stocks of Summer Steelhead that passed Bonneville Dam in 2019 during the Skamania and A-/B-Index Management Periods.

	Hatchery origin- No Clip	Subtotal for Skamania Period (Apr 1 - Jun 30)		Subtotal for A-/B-Index Period (Jul 01 - Oct 31)		Subtotal for A-/B-Index Period (Jul 01 - Oct 31)	
Reporting Group name	Reporting Group Code	Estimated abundance		A-INDEX: Estimated abundance		B-INDEX: Estimated abundance	
		Mean	95% CI	Mean	95% CI	Mean	95% CI

Lower Columbia	02_LOWCOL	0	0 – 0	0	0 – 0	0	0 – 0
Skamania	03_SKAMAN	0	0 – 0	0	0 – 0	0	0 – 0
Willamette	04_WILLAM	0	0 – 0	0	0 – 0	0	0 – 0
Big White Salmon	05_BWSALM	0	0 – 0	0	0 – 0	0	0 – 0
Klickitat	06_KLICKR	0	0 – 0	0	0 – 0	0	0 – 0
mid Columbia/Grande Ronde/Imnaha/lower Clearwater/lower Snake/lower Salmon	07_MGILCS	0	0 – 0	309	0 – 812	0	0 – 0
Yakima	08_YAKIMA	0	0 – 0	0	0 – 0	0	0 – 0
upper Columbia	09_UPPCOL	0	0 – 0	98	0 – 390	0	0 – 0
SF Clearwater	10_SFCLWR	0	0 – 0	1,120	627 – 1686	1,029	499 – 1640
upper Clearwater	11_UPCLWR	0	0 – 0	0	0 – 0	0	0 – 0
SF Salmon	12_SFSALM	0	0 – 0	0	0 – 0	0	0 – 0
MF Salmon	13_MFSALM	0	0 – 0	0	0 – 0	0	0 – 0
upper Salmon	14_UPSALM	0	0 – 0	193	0 – 578	0	0 – 0
Total		0		1,719		1,029	

Note: Based on the sample data described in Table 62 we estimated abundances of six categories of steelhead; i.e., clipped hatchery (H), unclipped hatchery (HNC), and natural-origin (W) for A-Index and B-Index fish. The sample was divided into the most strata possible maintaining a minimum sample of 100 clipped and unclipped steelhead per stratum in the A-/B-Index period. All weeks were pooled into a single stratum in the Skamania period.

Table 65. Estimated abundance of natural-origin unclipped stocks (excluding hatchery-origin unclipped) of Summer Steelhead that passed Bonneville Dam in 2019 during the Skamania and A-/B-Index Management Periods.

	Natural origin- No Clip	Subtotal for Skamania Period (Apr 1 - Jun 30)		Subtotal for A-/B-Index Period (Jul 01 - Oct 31)		Subtotal for A-/B-Index Period (Jul 01 - Oct 31)	
Reporting Group name	Reporting Group Code	Estimated abundance		A-INDEX: Estimated abundance		B-INDEX: Estimated abundance	
		Mean	95% CI	Mean	95% CI	Mean	95% CI
Lower Columbia	02_LOWCO L	234	0 – 468	0	0 – 0	119	0 – 358
Skamania	03_SKAMA N	585*	117 – 820	857	326 – 1476	119	0 – 358

Willamette	04_WILLAM	0	0 – 0	0	0 – 0	0	0 – 0
Big White Salmon	05_BWSALM	0	0 – 0	0	0 – 0	0	0 – 0
Klickitat	06_KLICKR	351	117 – 702	924	439 – 1538	0	0 – 0
mid Columbia/Grande Ronde/Imnaha/lower Clearwater/lower Snake/lower Salmon	07_MGILCS	351	117 – 702	19,996	18209 – 21593	214	0 – 478
Yakima	08_YAKIMA	0	0 – 0	1,390	731 – 2154	0	0 – 0
upper Columbia	09_UPPCOL	0	0 – 0	1,050	465 – 1668	0	0 – 0
SF Clearwater	10_SFCLWR	0	0 – 0	434	168 – 728	498	230 – 817
upper Clearwater	11_UPCLWR	0	0 – 0	206	54 – 418	0	0 – 0
SF Salmon	12_SFSALM	0	0 – 0	68	0 – 203	35	0 – 101
MF Salmon	13_MFSALM	0	0 – 0	704	281 – 1233	0	0 – 0
upper Salmon	14_UPSALM	117	0 – 351	5,314	4010 – 6628	68	0 – 203
	Total	1,639		30,943		1,054	

Note: Based on the sample data described in Table 62 we estimated abundances of six categories of steelhead; i.e., clipped hatchery (H), unclipped hatchery (HNC), and natural-origin (W) for A-Index and B-Index fish. The sample was divided into the most strata possible maintaining a minimum sample of 100 clipped and unclipped steelhead per stratum in the A-/B-Index period. All weeks were pooled into a single stratum in the Skamania period.
 *A small number (N=117) were estimated to be B-Index size.

Table 66. Estimated abundance of hatchery-origin clipped and unclipped broodstocks of A-/B-Index Summer Steelhead that passed Bonneville Dam in 2019 (July 1 – Oct 31).

Hatchery	Stock	GSI RepGrp	Broodstock	Brood year	Hatchery clipped				Hatchery unclipped			
					A-INDEX		B-INDEX		A-INDEX		B-INDEX	
					MLE	95% CI	MLE	95% CI	MLE	95% CI	MLE	95% CI

Skamania	Summer	03_SKA MAN	OmySKH1 6_su	2016	2,89 3	1732 – 3772	0	0 – 0	0	0 – 0	0	0 – 0
Round Butte	Deschutes River	07_MGI LCS	OmyRB17	2017	235	0 – 634	0	0 – 0	0	0 – 0	0	0 – 0
Umatilla	Minthorn Springs	07_MGI LCS	OmyUMA 17	2017	220	0 – 450	0	0 – 0	0	0 – 0	0	0 – 0
Round Butte	Deschutes River	07_MGI LCS	OmyRB16	2016	616	203 – 1119	0	0 – 0	98	0 – 293	0	0 – 0
Eastbank	Chelan/Methow/Okanoga n/Wenatchee	09_UPP COL	OmyEAST BK17	2017	0	0 – 0	0	0 – 0	98	0 – 293	0	0 – 0
Wells	On Station	09_UPP COL	OmyWEL 17	2017	118	0 – 355	0	0 – 0	0	0 – 0	0	0 – 0
Wells	On Station	09_UPP COL	OmyWEL 16	2016	670	225 – 1226	0	0 – 0	0	0 – 0	0	0 – 0
Winthrop NFH	Methow River	09_UPP COL	OmyWTP1 6	2016	112	0 – 336	0	0 – 0	0	0 – 0	0	0 – 0
Little Sheep Creek	-	07_MGI LCS	OmyLSCR 17S	2017	1,34 6	678 – 2067	0	0 – 0	0	0 – 0	0	0 – 0
Lyons Ferry	Grande Ronde	07_MGI LCS	OmyCGR W17S	2017	4,82 3	3533 – 6077	0	0 – 0	0	0 – 0	0	0 – 0
Lyons Ferry	Wallowa	07_MGI LCS	OmyWAL W17S	2017	1,83 0	1087 – 2681	0	0 – 0	0	0 – 0	0	0 – 0
Lyons Ferry	Tucannon	07_MGI LCS	OmyTUC W17S	2017	623	112 – 1196	0	0 – 0	86	0 – 259	0	0 – 0
Lyons Ferry	Touchet	07_MGI LCS	OmyTOU W17S	2017	0	0 – 0	0	0 – 0	27	0 – 81	0	0 – 0
Little Sheep Creek	-	07_MGI LCS	OmyLSCR 16S	2016	1,09 7	456 – 1864	0	0 – 0	0	0 – 0	0	0 – 0
Lyons Ferry	Grande Ronde	07_MGI LCS	OmyCGR W16S	2016	1,53 6	860 – 2264	0	0 – 0	0	0 – 0	0	0 – 0
Lyons Ferry	Wallowa	07_MGI LCS	OmyWAL W16S	2016	1,24 4	584 – 2002	0	0 – 0	0	0 – 0	0	0 – 0
Lyons Ferry	Tucannon	07_MGI LCS	OmyTUC W16S	2016	0	0 – 0	0	0 – 0	98	0 – 293	0	0 – 0
Dworshak NFH	Clearwater River	10_SFCL WR	OmyDWO C17S	2017	1,09 5	762 – 1479	0	0 – 0	280	146 – 439	0	0 – 0

Dworshak NFH	SF Clearwater	10_SFCL WR	OmySFC W17S	2017	306	148 – 503	0	0 – 0	81	27 – 162	0	0 – 0
Dworshak NFH	Upper Salmon	10_SFCL WR	OmyUSAL 17S	2017	0	0 – 0	0	0 – 0	35	0 – 105	0	0 – 0
Dworshak NFH	Upper Salmon	10_SFCL WR	OmyUSAL 16S	2016	506	230 – 806	211	57 – 387	429	153 – 721	0	0 – 0
Dworshak NFH	Clearwater River	10_SFCL WR	OmyDWO C16S	2016	1,876	1269 – 2420	3,818	3242 – 4452	295	115 – 505	1,029	685 – 1423
Oxbow	-	14_UPS ALM	OmyOXB O17S	2017	2,032	1035 – 3058	0	0 – 0	0	0 – 0	0	0 – 0
Pahsimeroi	Salmon River	14_UPS ALM	OmyPAH H17S	2017	2,044	1355 – 2813	0	0 – 0	0	0 – 0	0	0 – 0
Sawtooth	Salmon River	14_UPS ALM	OmySAW T17S	2017	2,953	2062 – 3833	0	0 – 0	0	0 – 0	0	0 – 0
Sawtooth	East Fork Salmon River	14_UPS ALM	OmyEFS W17S	2017	0	0 – 0	0	0 – 0	193	0 – 485	0	0 – 0
Oxbow	-	14_UPS ALM	OmyOXB O16S	2016	1,899	1067 – 2696	0	0 – 0	0	0 – 0	0	0 – 0
Pahsimeroi	Salmon River	14_UPS ALM	OmyPAH H16S	2016	1,209	662 – 1838	64	0 – 193	0	0 – 0	0	0 – 0
Sawtooth	Salmon River	14_UPS ALM	OmySAW T16S	2016	1,166	552 – 1881	0	0 – 0	0	0 – 0	0	0 – 0
#N/A	#N/A	#N/A	Unassigned	#N/A	1,180	608 – 2553	0	0 – 23	0	0 – 0	0	0 – 0
				TOTAL	33,628		4,093		1,719		1,029	

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Post-season analysis of Sockeye Salmon passing Bonneville Dam in 2019

We provided a timely post-season analysis of Sockeye Salmon once the majority of samples had been collected at the Bonneville Dam AFF in mid-August and distributed the report to USvOR TAC on August 9, 2019 (Table 47). In recent years, the results of our genetic analysis have become an official component of the post-season run reconstruction and pre-season forecasting that TAC performs in November of each year. Therefore, timely reporting of the individual assignments of the Bonneville Dam sample of Sockeye Salmon to GSI reporting groups is critical for efficient execution of fisheries management of this species in the Columbia River. In 2019, there were 1,857 Sockeye Salmon that were sampled at the AFF and genotyped for this analysis (Table 67). For the first year since we began genetic analysis of Sockeye salmon, we estimated zero abundance for the ESA listed stock (Redfish Lake Sockeye Salmon from the Snake River) (Table 67, Figure 37). The genetic analyses provide additional information that cannot be obtained by conventional tagging methods. For example, PIT-tags placed in adults at the Bonneville Dam AFF can identify the stock-of-origin only for fish that survive and are detected at upstream dams. However, the genetic analysis can provide stock-of-origin on most fish regardless of whether they survive further upstream of Bonneville Dam. This ability has allowed greater numbers of fish to be detected from the relatively rare ESA listed Snake River stock, which improves accuracy and precision of abundance estimates. In addition, the Lake Billy Chinook stock from the Deschutes River is rare and difficult to detect with PIT-tags because of limited arrays in the Deschutes River. Therefore, the ability for the genetic baseline to identify individuals from this stock provides the only way to effectively monitor this stock's abundance in the mainstem Columbia River. In 2019, we estimated the Lake Billy Chinook stock was 264 (95% C.I.: 143 – 409) fish. Similar to the 2018 year, we were able to estimate the reintroduced stock from the Yakima River using our PBT baseline; the stock abundance in 2019 was 265 (95% C.I.: 0 – 1103).

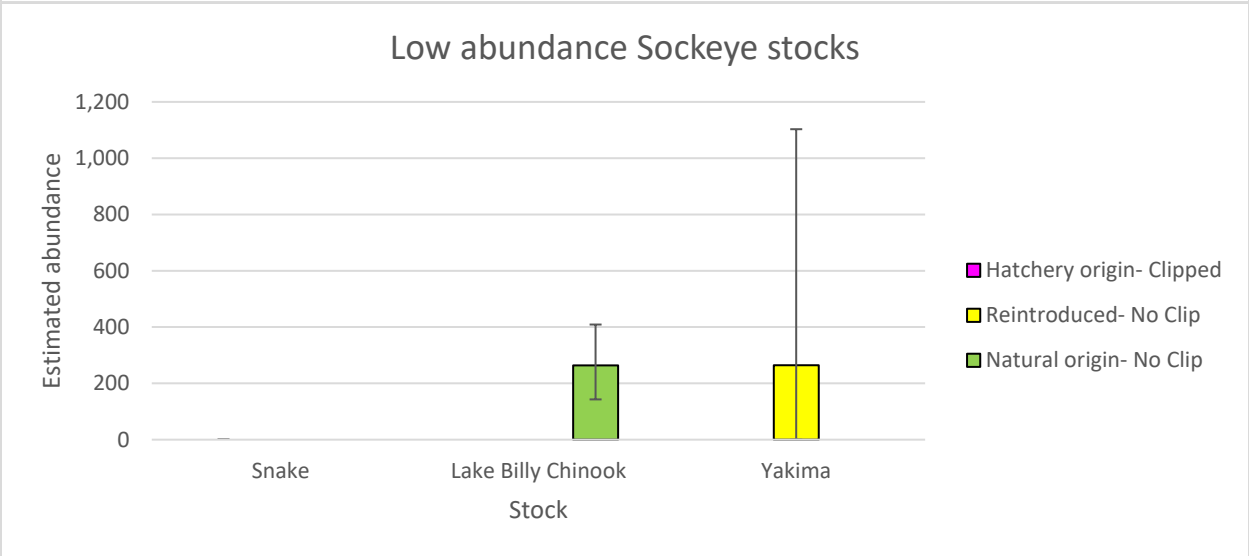
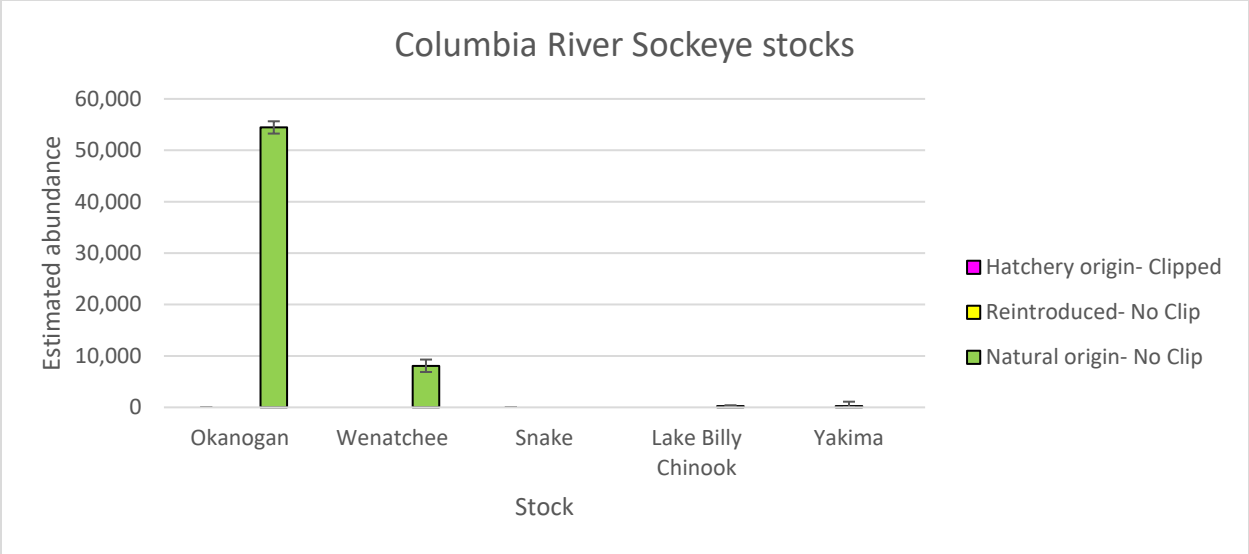


Figure 37. Abundance estimates of Sockeye Salmon stocks passing Bonneville Dam in 2019.

2056 **Table 67. Estimated abundance of Sockeye Salmon genetic stocks that passed Bonneville Dam in 2019.**

Reporting Group name	Hatchery origin- Clipped		Reintroduction- No Clip		Natural origin- No Clip	
	Estimated abundance		Estimated abundance		Estimated abundance	
	Mean	95% CI	Mean	95% CI	Mean	95% CI
Okanogan	0	0 – 0			54,466	53254 – 55645
Wenatchee					8,052	6878 – 9299
Snake	0	0 – 0				
Lake Billy Chinook					264	143 – 409
Yakima			265	0 – 1103		
Total	0		265		62,781	

2057 Note: The abundance is estimated from the total fish counts at the fish ladder windows at Bonneville Dam. Most stocks are identified
 2058 by GSI assignment, however this year we were able to use PBT to identify Yakima fish that were reintroduced.

Discussion

Parentage based tagging (PBT) and genetic stock identification (GSI) may be considered as methods that could replace the central functions of the coded wire tag program and could be a replacement for adipose fin marking to identify hatchery origin fish. However, this replacement would be contingent on continued genotyping of hatchery broodstock, fish passing Bonneville Dam, and harvested fish. For ocean fisheries management, additional hatcheries throughout the range of Chinook salmon would have to contribute broodstock samples to this PBT baseline in order for the method to serve ocean fisheries management and the need to monitor total fishery impacts for stocks including Columbia River stocks of fall Chinook (tules and upriver brights) harvested in ocean fisheries. The genetic methods provide a substantial amount of information when they are combined and used to analyze Columbia River Chinook salmon and steelhead passing Bonneville Dam. PBT improves the accuracy for defining hatchery-origin and by subtraction, total natural-origin stocks. Expansion of our PBT baseline to include hatcheries in the Columbia River has increased the proportion of hatchery origin fish passing Bonneville Dam that can be assigned to their broodstock source. We were able to assign the 78% of fish that were hatchery-origin summer A-/B-Index steelhead at Bonneville Dam in 2018 to 29 broodstock sources. As this effort continues to expand, we anticipate a corresponding increase in the proportion of hatchery origin fish that can be confidently assigned to their hatchery broodstock source. We have observed at least one sign of improved PBT coverage based upon our in-season analyses of Chinook Salmon that passed Bonneville Dam in 2019. Namely, the percentage of adipose-clipped fish abundance that was identified as PBT broodstock ranged from 85-100%, 94-100%, and 82-100% in the spring, summer, and fall management periods, respectively. We would expect these percentages to be high (~100%) only if all hatchery broodstock above Bonneville Dam are included in the PBT baseline AND our estimated tag rates are accurate for each hatchery broodstock. The fact that the management periods mostly attained high percentages gives us confidence that the PBT baseline coverage is nearly complete for Chinook Salmon above Bonneville Dam.

Genetic monitoring combining PBT and GSI is one of a number of possible tools that can be used to identify hatchery and natural fish at various resolutions. Other methods include, CWTs, PIT tags, VIE tags, and otolith marks. Adipose fin clips can be used to differentiate hatchery fish from wild fish either when fish are clipped at 100% or through expansions if stocks are not clipped at 100%. PBT can further discriminate among hatchery stocks within the reporting groups that we use for GSI analyses, and so we can now characterize different age-classes from particular hatcheries by run-timing distributions and estimate their abundance at Bonneville Dam. GSI continues to provide information that would not be possible with PBT, especially for natural-origin stocks.

This long-term study will allow us to characterize trends in run timing and abundance of steelhead and Chinook and Sockeye salmon and provide this data to fisheries managers. We were able to address the following **F&W Program Management Questions:**

What are the status and trend of adult productivity of fish populations?

What are your in-river monitoring results and what are your estimates of stock composition and stock-specific abundance, escapement, catch, and age distribution?

Trapping at Bonneville Dam can only be done at very low rates due to restrictions placed on trap operations by USACE and NFMS. Low sample rates inhibit getting a representative

sample of various stocks of fish. Higher sample rates would improve the precision of the estimates of fish at Bonneville Dam. Some fisheries were also sampled at very low rates.

We identified 13 Chinook salmon hatchery-origin stocks (66 hatchery broodstock sources) and 7 natural-origin stocks estimated to have relative abundances $\geq 1,000$ fish passing Bonneville Dam in 2018. The migratory delays observed at Bonneville Dam in the spring of 2018 appear to have affected the run-timing distributions for several hatchery-origin and natural-origin stocks. These delays likely contributed to the broad overlap in the temporal distributions of spring, summer/fall, and fall Chinook stocks, and our detection of spring stocks passing Bonneville Dam well into the summer management period.

We identified five steelhead hatchery-origin stocks (29 hatchery broodstock sources) and five natural-origin stocks estimated to have relative abundances $\geq 1,000$ fish passing Bonneville Dam in 2018. We found that genetic stocks seemed to fit well into the historical management categories, particularly the hatchery-origin stocks. B-Index fish appeared to have later run-timing compared to A-Index fish from the same stock.

Characteristics of steelhead that assigned to Snake River hatchery broodstock sources generally support the typical A-run and B-run steelhead life history categories. The relatively large (≥ 78 cm) steelhead were found primarily to originate from Dworshak hatchery broodstock. These fish were also relatively old (2- and 3-ocean-age) and were derived from the Clearwater R., which is one of the regions expected to produce “B-run” steelhead. It is notable that the MGILCS reporting group represents some fish both within and outside the Snake River steelhead DPS, but does not represent all of the fish within the Snake River DPS.

This was the seventh year that we were able to analyze sockeye salmon using GSI, and similar to analyses for 2017, in 2018, we used a GSI baseline that included kokanee samples from the Deschutes River drainage, Wallowa Lake, and other locations on the Snake River. We estimated relative stock composition and stock abundance for sockeye passing Bonneville Dam in 2018, and found that the Okanogan stock has the greatest relative abundance followed by the Wenatchee stock. We found fewer fish from the Snake River stock compared to previous years, and identified fish from Lake Billy Chinook and even from the reintroduced stock in Yakima River using a PBT baseline. We also found that the migratory run timing for the Okanogan and Wenatchee stocks overlaps broadly at Bonneville Dam, with the Snake stock migrating slightly later in the year.

This year (2019) we continued the increased frequency of in-season reporting as we performed in 2018 during the Chinook Salmon spring and summer management periods, and we have offered timely post-season reports for the fall management period of Chinook Salmon, as well as Skamania and A-/B-Index Summer Steelhead Management Periods, and Sockeye Salmon. For steelhead and sockeye, our genetic analyses have become officially integrated into the routine analyses that U.S. v OR TAC performs. In all cases, the genetic analyses are a complement to the data that fisheries managers have available to gain insights into the biological characteristics of fisheries. The genetic analysis of the Spring Chinook Salmon test fishery data may begin to play a particularly important role in the in-season analyses because of its potential predictive power for stock abundance and timing at Bonneville Dam weeks in advance of the arrival of the run.

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2160 [111Steele2010%20Parentage%20Based%20Tagging%20Snake%20River%20Steelhead%20Sal](https://research.idfg.idaho.gov/Fisheries%20Research%20Reports/Res11-111Steele2010%20Parentage%20Based%20Tagging%20Snake%20River%20Steelhead%20Salmon.pdf)
2161 [mon.pdf](https://research.idfg.idaho.gov/Fisheries%20Research%20Reports/Res11-111Steele2010%20Parentage%20Based%20Tagging%20Snake%20River%20Steelhead%20Salmon.pdf)

2162

2163 Section 5: Local adaptation in salmonids

2164 Introduction

2165 Environmental and landscape features can greatly contribute to the population structure,
2166 life history diversification, and local adaptation of organisms in aquatic habitats
2167 (reviewed in Storfer et al. 2006). Geographic barriers to dispersal include recent events
2168 that may have been human induced (e.g., dams) as well as ancient events such as
2169 glaciations and formation of mountain chains (e.g., Castric et al. 2001). However, other
2170 environmental characteristics such as elevation, temperature, forest cover, and
2171 precipitation may influence distribution, adaptation, and gene flow of species (Dionne et
2172 al. 2008; Narum et al. 2008). For example, the geographic distributions of species
2173 ranges' are often determined by thermal tolerance (Brannon et al. 2004) and may
2174 necessitate adaptations for survival in extreme environments (Rodnick et al. 2004).

2175
2176 Screening with many genetic markers provides the opportunity to investigate local
2177 adaptation in natural populations and identify candidate genes under selection (Beaumont
2178 and Nichols 1996; Beaumont and Balding 2004; Excoffier et al. 2009). This has become
2179 a commonly employed approach in ecological and population genetics studies to detect
2180 outlier loci that are putatively under selection (e.g., Vasemagi and Primmer 2005; Nosil
2181 et al. 2008). Additionally, correlation methods can be highly informative to identify
2182 markers in coding and cis-regulatory regions of known functional genes that are
2183 associated with specific selective pressures or phenotypes (Lyman and Mackay 1998;
2184 Chase et al. 2009; Torgerson et al. 2009). With increasing genomic information available
2185 for non-model organisms, single nucleotide polymorphisms (SNPs) have begun to see
2186 increased use as genetic markers for population genetic studies (e.g., Morin et al. 2004).
2187 These sequence polymorphisms are densely scattered throughout the genome of most
2188 organisms, and are commonly observed in both coding and non-coding regions of
2189 functional genes making them ideal markers to study adaptive molecular variation (e.g.,
2190 Akey et al. 2002). In a large suite of SNPs that are distributed across the genome (e.g.,
2191 Narum et al. 2018), it is possible to utilize both functionally neutral and adaptive markers
2192 within a single study. This combination of information provides a powerful approach to
2193 study questions in ecological genetics since both demographic processes (i.e., gene flow
2194 and genetic drift) and local adaptation (i.e., selection) may be inferred.

2195
2196 Molecular techniques such as RNA-seq (Wolf 2013) also provide the opportunity to
2197 investigate transcriptional response to thermal stress and further identify mechanisms for
2198 thermal adaptation. Patterns of gene expression under heat stress are important to
2199 determining evolutionary adaptation among conspecific populations that occupy various
2200 environments. Multiple genes have been shown to be involved in heat tolerance across
2201 many species, including highly conserved heat shock proteins (hsps) that are upregulated
2202 under stressful conditions such as exposure to heat (Morimoto et al. 1992; Sorensen et al.
2203 2003). An adaptive heat shock response has additionally been shown to occur among
2204 conspecific populations that occupy variable environments (e.g., Dahlhoff and Rank
2205 2000; Sorensen et al. 2001). However, many genes are known to have a role in
2206 regulating the effects of temperature and are likely to be involved in thermal adaptation

(Sorensen et al. 2005; Kassahn et al. 2007). Thus, RNA-seq provides the opportunity to investigate differential expression across the transcriptome and identify biological pathways involved in evolutionary response to thermal stress.

Thus, genome scans with large numbers of SNP markers (e.g., RAD sequencing, Baird et al. 2008; Pool-seq, Schlotterer et al. 2014) and gene expression (e.g., RNA-seq) approaches may be effective tools for identifying the genetic architecture underlying specific traits such as thermal tolerance, run-timing/maturation, disease resistance, anadromy, and age-at-maturity. Once these underlying genomic regions are identified, they can be broadly screened in populations throughout the Columbia River Basin to facilitate management for long term conservation and recovery of salmonids.

Fish Population RM&E

F&W Program Strategy: Assess the status and trend of diversity of natural and hatchery origin fish populations.

F&W Program Management Question: What are the status and trend of diversity of natural and hatchery origin fish populations?

Uncertainty Research

See Appendix A for table of critical uncertainties research.

Project Map:

<http://www.cbfish.org/Project.mvc/Map/2009-005-00>

Contract Map(s):

<http://www.cbfish.org/Contract.mvc/Map/61839>

<http://www.cbfish.org/Contract.mvc/Map/65575>

Methods: Protocols, Study Designs, and Study Area

Method Title: Whole Genome Resequencing

- **Method Link:**

- [Whole Genome Resequencing: Poolseq Individual Barcode v1.0](#) (ID: 6754) Published

- [Whole Genome Resequencing: Poolseq Pooled v1.0](#) (ID: 6775) Published

Method Summary:

With reference genome assemblies now publicly available for both Chinook salmon and steelhead, this enables whole genome resequencing for investigating adaptive variation across a large portion of the genome (50-80%) in these species. Whole genome resequencing methods include individually barcoded samples, or pools of samples (Pool-seq; Schlotterer et al. 2014) depending on the study design. For both methods, sequence data is aligned to the reference genome assembly, and allele frequencies from millions of SNPs are analyzed to detect statistically significant regions of the genome associated with

specific traits or adaptation to environmental factors. Putatively neutral regions of the genome are also useful for standard phylogeny and demographic analyses of populations. In most studies, allele frequencies are available for collections but sequencing depth is typically not high enough to provide individual genotypes. However, candidate SNPs may be developed into standard panels with GTseq or other approaches to genotype many individuals to validate trait association, determine inheritance, and estimate linkage disequilibrium.

Method Title: RAD sequencing v1.0

Method Link: <http://www.monitoringmethods.org/Method/Details/4144>

Method Summary:

RAD sequencing is a technique for tagging DNA at restriction enzyme cut sites with adapters used in massively parallel sequencing. This method allows thousands of SNPs to be discovered and genotyped in several individuals. Through the use of sample specific DNA barcodes included in the adapters, information for specific samples can be separated in silico following sequencing. This method effectively reduces sequence complexity by targeting only sequence surrounding restriction enzyme cut sites making alignments among sequencing reads far less computationally intense. The sequence alignments among samples can then be analyzed for both identification and genotyping of SNPs (Single Nucleotide Polymorphisms). This method was first described by Baird et al. (2008).

Method Title: Obtain gene expression data via RNAseq v1.0

Method Link: <http://www.monitoringmethods.org/Method/Details/607>

Method Summary:

Compare gene expression between fish of different genetic backgrounds but raised in the same environment. Molecular techniques such as RNAseq provide the opportunity to investigate transcriptional response and further identify mechanisms for thermal adaptation. Patterns of gene expression are important to determining evolutionary adaptation among conspecific populations that occupy various environments.

Results

Objective 1) Environment & Landscape Genetics

Hypotheses:

Hypothesis 1: Environmental and landscape features act as drivers of selection leading to local adaptation of fish populations. Testing many variables can identify the key environmental drivers of selection.

Hypothesis 2: Environmental drivers of selection act on specific genes resulting in different allelic and genotypic frequencies among adapted populations. Genome scans can identify candidate genes involved in local adaptation of fish populations across heterogeneous landscapes.

Activities implemented:

Multiple studies have been initiated to investigate local adaptation with a landscape genetics/genomics approach to address the two hypotheses above. Our work has focused on steelhead and Chinook salmon throughout the Columbia River Basin. Studies continue to progress as more markers become available throughout the genome of each species.

For steelhead, landscape genetics work has been done at both fine and broad scales. Fine scale landscape genetics approaches were used to identify ecological patterns of residence vs anadromy and found consistent evidence that certain landscape features have led to prevalence of certain life history types in *O. mykiss* (Narum et al. 2008a) but also *O. nerka* (Nichols et al. 2016). Broad scale studies were initially done with panel of 188 SNP markers to investigate patterns of landscape genetics across 145 populations in the Columbia River Basin (Matala et al. 2014). Results indicated that precipitation and temperature were the primary environmental drivers of local adaptation and neutral genetic structure largely reflected isolation by distance. These results were shown to be robust due to replication of multiple populations representing each distinct genetic unit (Hand et al. 2016). A follow-up study (Micheletti et al. 2018a) was done with a much larger number of genetic markers (~20K SNPs) at broad scale and found evidence that the mainstem Columbia River migratory corridor exhibits greater selective pressure on steelhead than natal tributaries (Figure 38). Populations that must migrate long distances through the migratory corridor were under the most intense selection and multiple candidate genes were identified (Micheletti et al. 2018a). Additional studies are underway to further pinpoint candidate genes through whole genome resequencing.

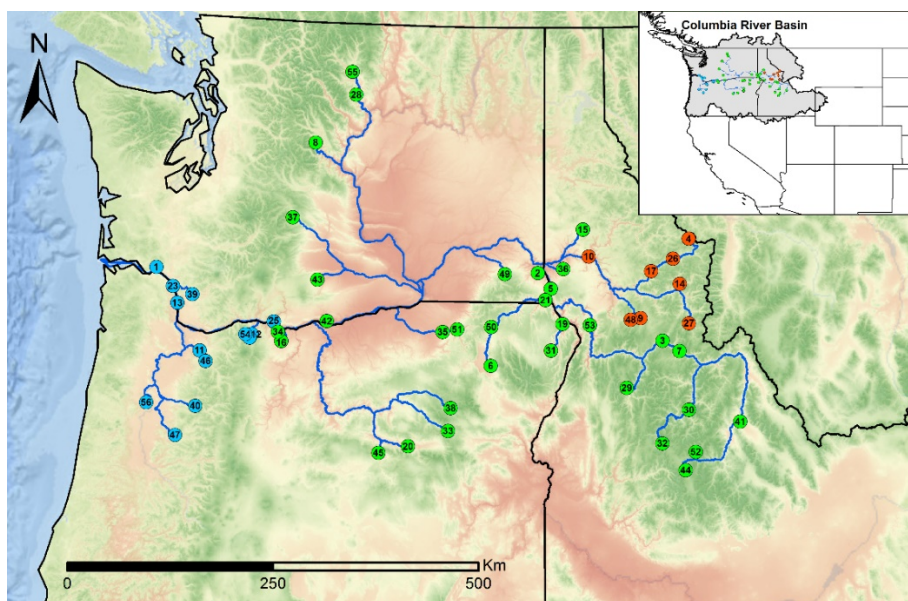


Figure 38. Steelhead collection localities colored by genetic cluster (from Micheletti et al. 2018a). blue = coastal, green = inland, red = inland-Clearwater. Blue lines indicate each population's migration path to the ocean. Map layer shows mean annual temperature for the warmest quarter with transition from colder (green) to warmer (red).

For Chinook salmon, landscape genetics work has also been done at both fine and broad scales. Initial studies evaluated the utility of certain marker types for landscape genetics and found consistency in signals among marker types (Narum et al. 2008b; Hess et al. 2011), but that SNPs provided greater potential for identifying candidate genes involved in local adaptation and should be used in subsequent studies (Narum et al. 2013a). Broad scale studies were initially done with panel of 96 SNP markers to investigate patterns of connectivity related to landscape features across 54 populations in the Columbia River Basin (Matala et al. 2011). Results indicated that precipitation, elevation, and temperature were the primary environmental drivers of local adaptation depending on genetic lineage, and neutral genetic structure largely reflected isolation by distance within each lineage (Matala et al 2011). A follow-up study (Hecht et al. 2015) was done with a much larger number of genetic markers (~20K SNPs) at broad scale and found that between 6-22% of genetic variation could be accounted for by environmental features such as precipitation, temperature, and migration distance (**Figure 39a**). Several candidate markers were associated with local adaptation within and among lineages (**Figure 39b**; Hecht et al. 2015). A follow-up study to investigate local adaptation through whole genome resequencing has demonstrated that a broad portfolio of diversity persists in Chinook salmon and candidate genes for local adaptation are widespread throughout the genome within and among lineages (Narum et al. 2018). Additional studies are underway to represent genome wide variation for a more thorough set of populations throughout the Columbia River Basin.

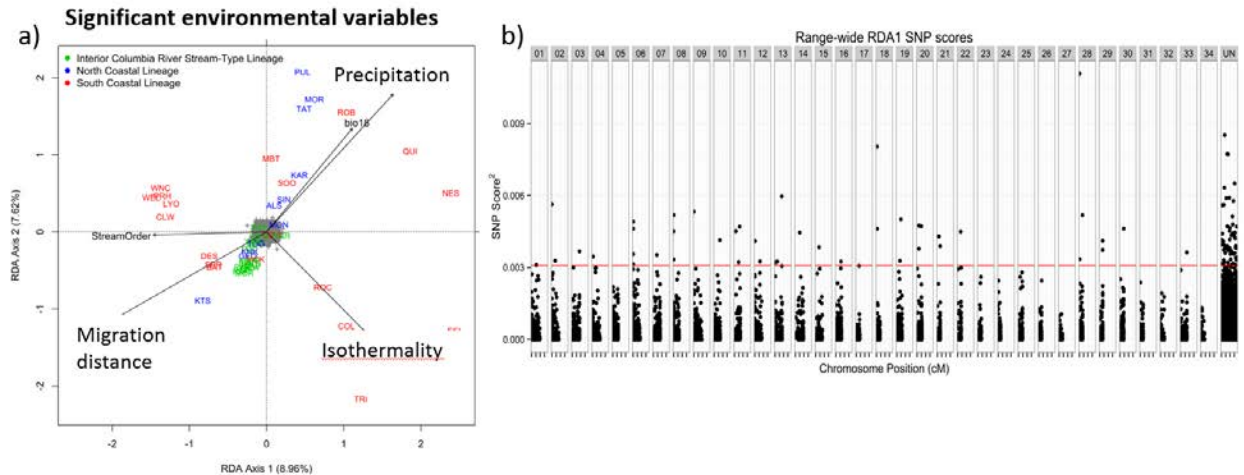


Figure 39. a) Environmental drivers of selection and b) candidate markers for local adaptation in Chinook salmon (from Hecht et al. 2015). a) Population scores for canonical RDA axis 1 and 2 are represented by the three-letter abbreviation for each population, coloured to represent the lineage assignment of that population. SNP variance is indicated by the position of grey '+' symbols radiating from the plot centroid, each representing a different locus. Environmental factors are depicted as black vectors (arrows), where the length of the vector is a representation of the magnitude of the contribution of that environmental variable in explaining SNP variance. The angle between environmental variable vectors is a representation of the correlation between those variables. Vectors and points are plotted with symmetrical scaling (scale = 3) to preserve the relationship between scores, without focusing on a single score.

Summary of accomplishments & lessons learned:

Steady progress has been made to better understand landscape genomics and neutral vs. adaptive patterns of genomic variation for steelhead and Chinook salmon as reflected in 11 publications (Narum et al. 2008a; Narum et al. 2008b; Matala et al. 2011; Hess et al. 2011; Narum et al. 2013a; Matala et al. 2014; Hecht et al. 2015; Hand et al. 2016; Nichols et al. 2016; Micheletti et al. 2018a; Narum et al. 2018).

Results consistently show specific environmental variables are drivers of selection in both steelhead and Chinook including precipitation, temperature, and migration distance through the mainstem Columbia River. Neutral variation across studies consistently accounts for highly distinct lineages of each species, and genetic structure within each lineage that is related to geographic location by sub-basins. While adaptive genetic variation can be evident with modest numbers of SNP markers, specific candidate genes are best pinpointed by genome resequencing.

Recent work has focused on collecting genome resequencing data for several populations of Chinook salmon and steelhead to enable landscape genomics analyses with high density markers throughout the genome. This data overlaps with collections that were sequenced under Section 2 (Table 3 & Table 5) that will also be used for baseline allele frequencies. Landscape genomics analyses for both Chinook salmon and steelhead are ongoing.

Objective 2) Genetic Basis for Phenotypic Expression of Traits

Hypotheses:

Hypothesis 1: Many phenotypic traits include a heritable component that must have a genetic basis, which can be detected with association mapping and gene expression to identify candidate genes associated with specific traits.

Hypothesis 2: Markers from candidate genes can be developed into cost efficient assays for genotyping large numbers of individuals to monitor genetic variation for phenotypic traits at broad scales, with the potential to predict resiliency of populations to environmental changes.

Activities implemented:

Multiple studies have been initiated to investigate the genetic basis for multiple phenotypic traits using association mapping or gene expression approaches. Work has focused on genomic regions associated with several different traits. Work on resident vs. anadromous *O. mykiss* (Narum et al. 2008a; Hecht et al. 2013) has revealed that a combination of environmental and genetic factors contribute to determination of these life history types (Figure 43).

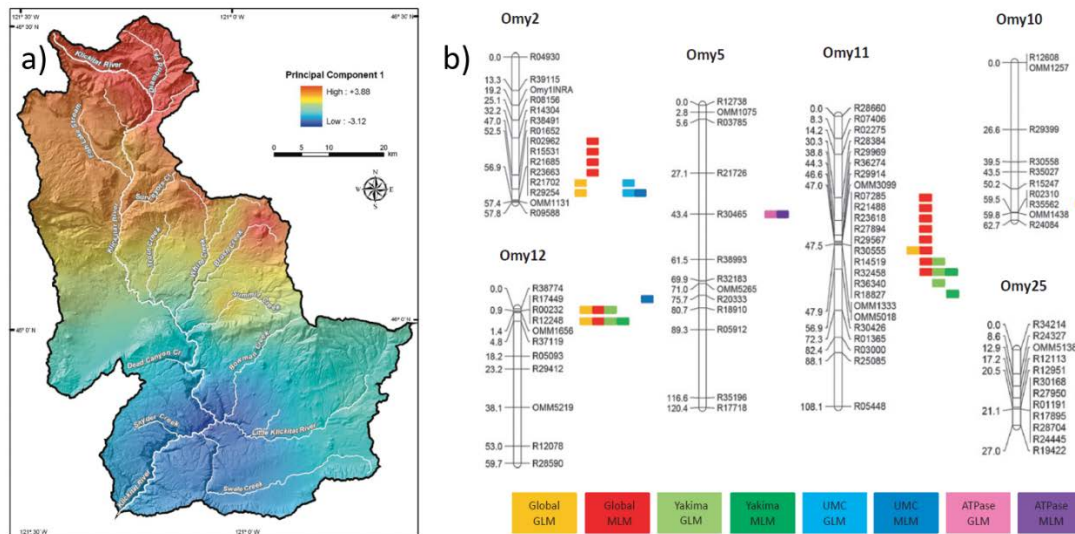


Figure 40. a) Synthesis map illustrating interpolation principal component (PC1) that accounts for 59% of variation in resident (red) vs anadromous (red) life history (from Narum et al. 2008a). b) Significant loci detected from eight individual association tests, where each test is represented by a different colored pill and placed to the right of the linkage group. GLM, general linear model; MLM, mixed linear model (from Hecht et al. 2013).

Run-timing related to maturity in steelhead (Hess et al. 2016; Micheletti et al. 2018c) and Chinook salmon (Hess and Narum 2011; Narum et al. 2018) has been shown to be controlled by a genomic region of major effect with the same candidate genes in both species (*GREB1L*, *ROCK1*, intergenic region; Figure 41).

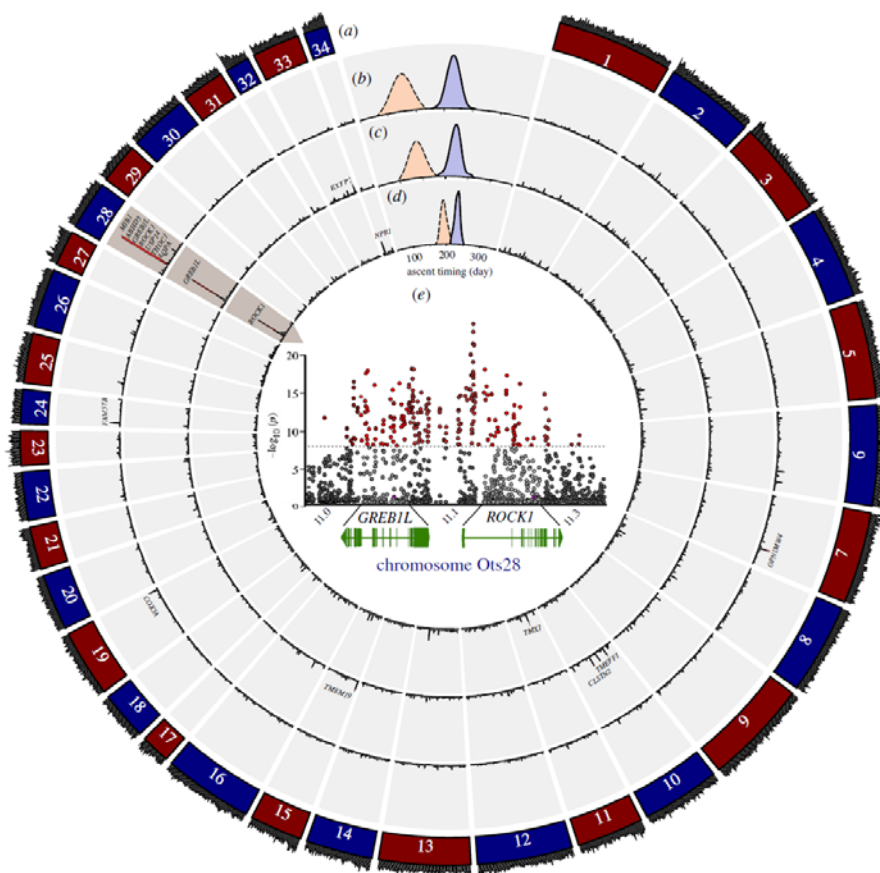


Figure 41. Manhattan plot for premature and mature collections of Chinook salmon (from Narum et al. 2018). (a) Sequence coverage (black outer ring) for each chromosome. (b) Significant divergence between premature (spring-run) and mature (fall-run) migrating Chinook salmon in the Cowlitz River of the coastal lineage. (c) Significant divergence between premature (Methow River summer-run) and mature (Priest Rapids fall-run) migrating Chinook salmon within the interior ocean-type lineage. (d) Significant divergence for Chinook salmon returning to Johnson Creek (interior stream-type) that enter freshwater premature (spring/summer-run), but the final ascent to spawning grounds is bimodal with early premature and late mature females. (e) Annotation of the 203 Kb region on Ots28 between 11.022 and 11.225 Mb (*GREB1L*, *ROCK1*, and intergenic regions) with significance based on CMH tests. Significant genes are labelled. Timing of ascent to spawning grounds for premature (dashed line, orange) and mature (solid line, blue) collection pairs are shown within the ring for each lineage (b–d). Purple dots show two non-synonymous SNPs.

Pooled-sequencing and a Cochran-Mantel-Haenszel (CMH; Mantel 1963) test executed by Micheletti et al. (2018c) detected steelhead markers associated with migration-timing on chromosome 28 within *GREB1L* and *ROCK1* genes and intergenic region between the two genes. With these markers identified, we can evaluate haplotype variation associated with migration-timing among steelhead populations collected across the Columbia River basin.

A principal component analysis (PCA) of allele frequencies of putatively neutral markers, without linkage disequilibrium (LD), will be plotted for all populations. Putatively neutral markers will be assessed to detect underlying population structure, which we expect to coincide with coastal and inland lineages. The PCA of putatively neutral markers will accompany a discriminate analysis of principal components (DAPC) with the R package adegenet 2.1.0 to assign probability of individual membership to genetic

groups, K , revealed with the putative neutral marker PCA (Jombart 2008, Jombart and Ahmed 2011). The DAPC recovers maximum genetic variation between groups, while minimizing genetic variation within groups (Jombart, 2008; Jombart and Ahmed, 2011). The ‘find.clusters’ adegenet function will run for 25 instances for $K=1$ through $K=10$. The Bayesian information criterion (BIC) will be averaged and scaled by the standard deviation for each K value. The most appropriate number of genetic groups will be determined with the greatest ΔK value as described in Evanno et al. (2005).

A total of 13 markers will be evaluated on chromosome 28 within the *GREBIL*, *rock1*, and intergenic region between *GREBIL* and *ROCK1* (Hess et al. 2016, Micheletti et al. 2018a). Premature, mature, and heterozygote genotypes migration-timing will be established based on genotypes from previous studies and the Skamania stock genotype, which is a hatchery-strain intensively selected for premature migration and cultured since 1956 with steelhead from the Washougal and Klickitat Rivers (Chilcote et al. 1986). Premature, mature, and heterozygote migration timing genotype proportions will be assessed across all collection locations. A PCA of allele frequencies of adaptive markers will also be conducted for all collection locations to assess genetic grouping based on migration-timing.

We seek to identify subsets of the markers, or haplotype blocks, most associated with migration-timing. Additionally, we will assess linkage disequilibrium (LD) within the 13 markers. Haplotype blocks of the 13 chromosome 28 markers will be defined with solid spine LD analysis in the Java Runtime Environment software, Haploview 4.2, across all collection locations (Barrett et al. 2005). The same markers will be assessed for LD in individuals from coastal and inland lineages (as delineated by DAPC) separately. The number and frequency of haplotypes across the basin will be compared between haplotype blocks. Variation of genotype proportions will also be evaluated with various sets of adaptive, chromosome 28 markers.

Redundancy analyses (RDA) will be conducted for all Columbia River basin sites to model the degree to which the variation in environmental variables explains the variation in allele frequencies of adaptive markers included in the haplotype blocks (Borcard et al. 1992; Kierepka and Latch 2015). Vegan 2.5-6 R package (Oksanen et al. 2019) will be used to analyze all populations together and each lineage separately. We will use environmental variables significantly associated with adaptive genetic variation in a previous study (Micheletti et al. 2018b). When two highly correlated (>0.75 pairwise correlation; Asuero et al. 2006) environmental variables are identified, one will be removed from further analyses and the variable kept will be determined from biological relevance to salmonids according to previous studies (Hecht et al. 2015; Olsen et al. 2011; Micheletti et al. 2018b). One-way analysis of variance (ANOVA) with a Tukey’s range test (Tukey 1949) will identify significant variability in salmonid habitat. Environmental variables will be analyzed with the “envfit” PCA function of the vegan R package. The ANOVA test and PCA together determine significant environmental variables within and among *O. mykiss* habitats that will be measured in this study. The final RDAs will be run with significant ($\alpha=0.05$) environmental variables retained from permutation tests with 1000 permutations. Frequency of alleles in the haplotype block associated with migration-timing are correlated to environmental variables with RDA

constraint scores. Constraint scores indicate the degree of correlation and whether there is a positive or negative relationship between environmental variables and allelic frequencies.

We are also analyzing a large sample of steelhead collected at Bonneville Adult Fish Facility (N=1,538) and in the Hood River (N=354) for patterns of association of the 13 chromosome 28 (*GREB1/ROCK1*) markers with two attributes of run-timing: the day each fish passed Bonneville dam and the day each fish was recorded at their most upstream passive integrated transponder (PIT) array (Figure 42). These samples were filtered from a larger set of complete PIT histories using custom R code, and consist of those samples for which the arrival to spawning tributary could be reliably inferred. Filtering was designed to exclude iteroparous spawning fish (kelts) with incomplete histories, “dip-in” fish that seek temporary thermal refuge in rivers other than their spawning destination, “overshoots” that travel farther upstream than necessary in higher order rivers but later regress, as well as fish only recorded at sites from which precision in arrival time could not be expected (e.g. dams and arrays in higher order rivers). While the Bonneville fish include mostly Interior Lineage fish, and are thus almost exclusively “summer-run” fish, the Hood River data includes both canonical “summer-run” and “winter-run” fish.

For Bonneville samples, in addition to the two raw measures of run time, we will calculate two compound measures, the lag time between Bonneville passage day and upstream arrival day, and the effective speed, which is the former compound statistic divided by the river distance between Bonneville Dam and the respective PIT array. Finally, we will also examine several relative measures: the Bonneville passage day or upstream arrival day relative to the median or last of such day within a sub-basin or hydrological unit, both within and across measures. In all, we will examine genetic association with 20 total measures of run timing.

From both sets of steelhead samples, we will infer patterns of linkage and haplotype frequencies using Haploview (Barrett, Fry, Maller, & Daly, 2005), which implements an expectation-maximization algorithm, and Shapeit, which utilizes a hidden-markov model and graph based algorithm (Delaneau, Marchini, & Zagury, 2012). Genetic association in these data will utilize both SNP-based and haplotype-based association analyses, including both the “mixed-linear” and “BLINK” models of GAPIT (Lipka et al., 2012) and the GLM-based haplotype score permutation routine of Haplo.stats (Sinnwell P & Schaid J, 2005). We expect these analyses to indicate 1) if the association of run timing with chromosome 28 markers is consistent between Coastal Lineage and Interior Lineage steelhead, 2) if the degree and pattern of correlation across the candidate region is the same, 3) whether heterozygotes for chromosome 28 markers have a phenotypic indicative of additive or dominant expression patterns, and 4) which markers or haplotypes are most strongly associated with run timing and may be most predictive of run timing from

genetic surveys, not unlike the results from Chinook salmon, discussed below.

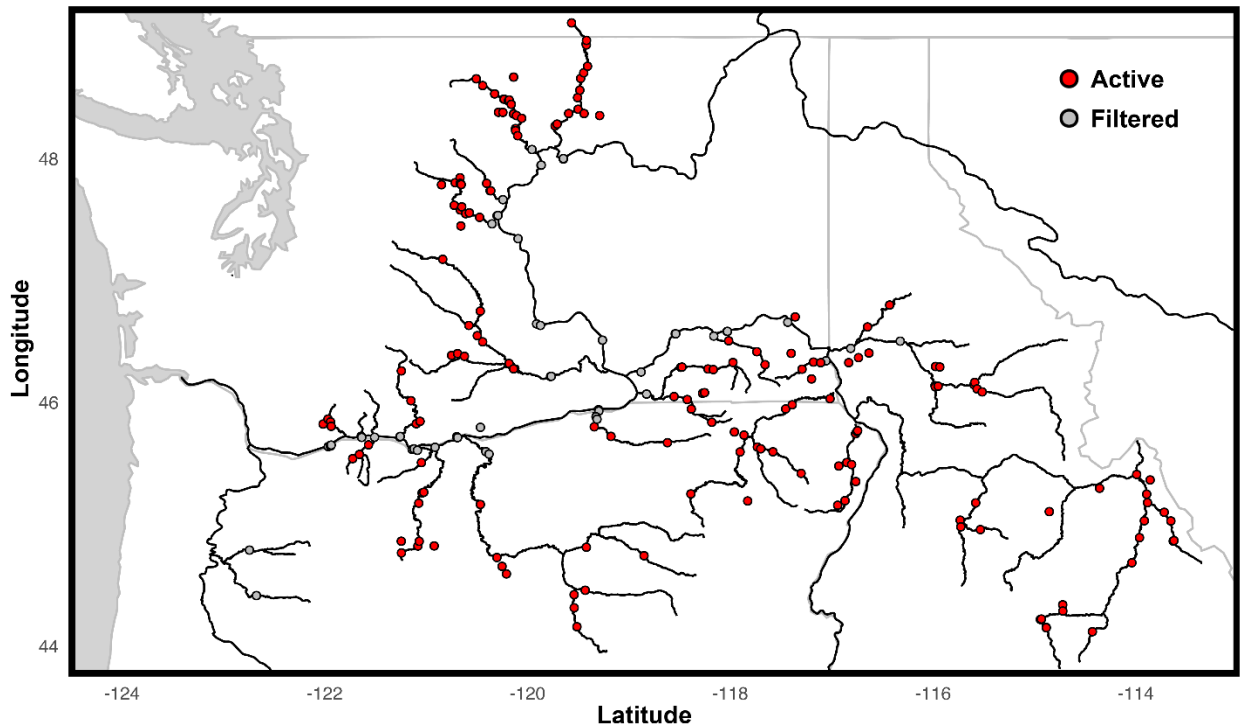


Figure 42. Map of passive integrated transponder (PIT) arrays at which steelhead passing and tagged at Bonneville Dam in 2013 to 2018 were later recorded. Filtered sites are those for which precise upstream arrival time could not be ensured, while fish recorded at the remaining active sites (N=1,538 samples) provide run-timing phenotype data for genetic association analyses.

For Chinook salmon, 33 candidate markers were developed that span 220kb on chromosome 28 including candidate genes *greb1L* and *rock1* for adult migration timing. Using individual-level genotypes from these candidate markers, we tested for an association with migration phenotypes across three distinct lineages that demonstrate both an early and late migration phenotype. We then used pedigree data from one of the populations, which enabled association tests between the candidate markers and fitness. Estimates of fitness were based on previous estimates of reproductive success from Janowitz-Koch et al. (2019).

A series of PCA results demonstrated clear differences in clustering between putatively neutral markers and those associated with adult migration timing in both the Lower Columbia and Interior ocean-type lineages (Figure 43). The pattern was investigated separately for the larger set of individual samples from the Interior stream-type lineage that had continuous data for adult migration timing including those fish that were intermediate between early and late migration timing peaks. There was substantial overlap between early and late migrating samples for the Interior stream-type lineage (Figure 43).

Association tests validated that the majority of markers were significantly associated with migration timing for all three lineages (Figure 44). The strongest association was consistently observed for markers within or upstream of the *rock1* gene, closely followed

by markers located within or upstream of greb1L. Out of the 33 candidate SNPs, 13 SNPs were significantly associated with fitness (

Figure 45).

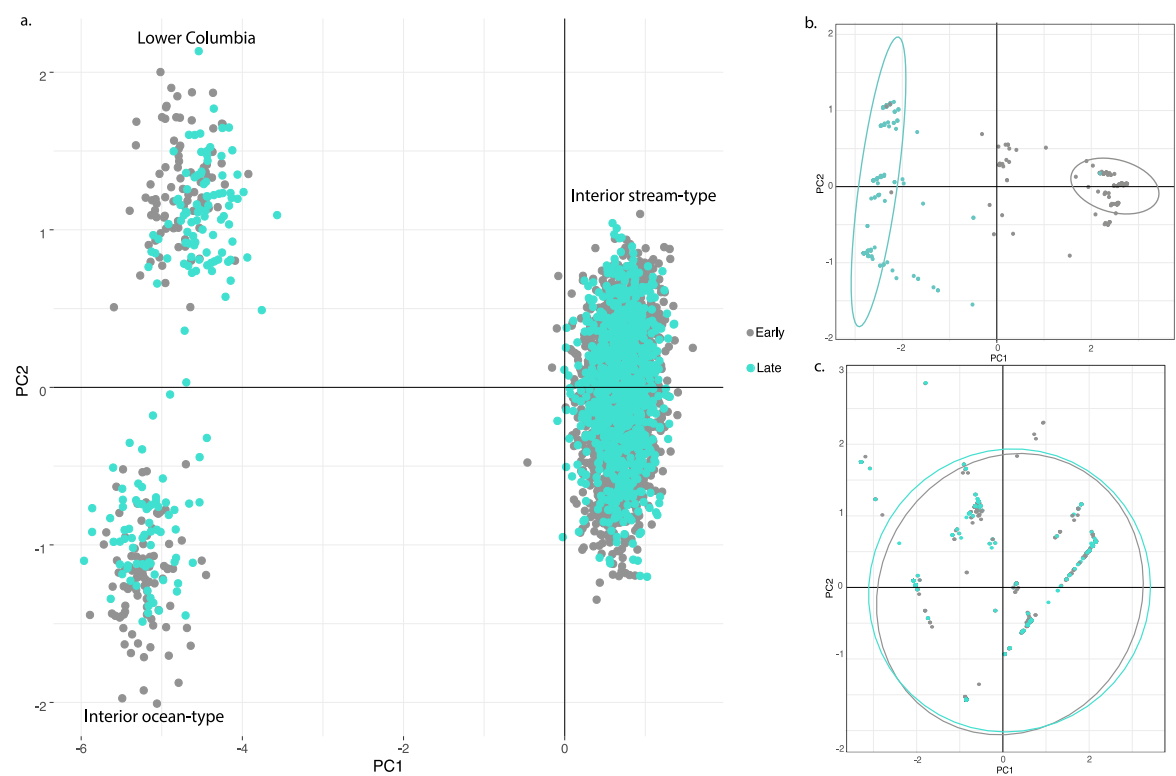


Figure 43. PCA of genetic variation in Chinook Salmon. Results presented represent A) 185 neutral SNP markers, B) 33 chromosome 28 markers for the Lower Columbia and Interior ocean-type populations combined, and C) 33 chromosome 28 markers for the Interior stream-type population.

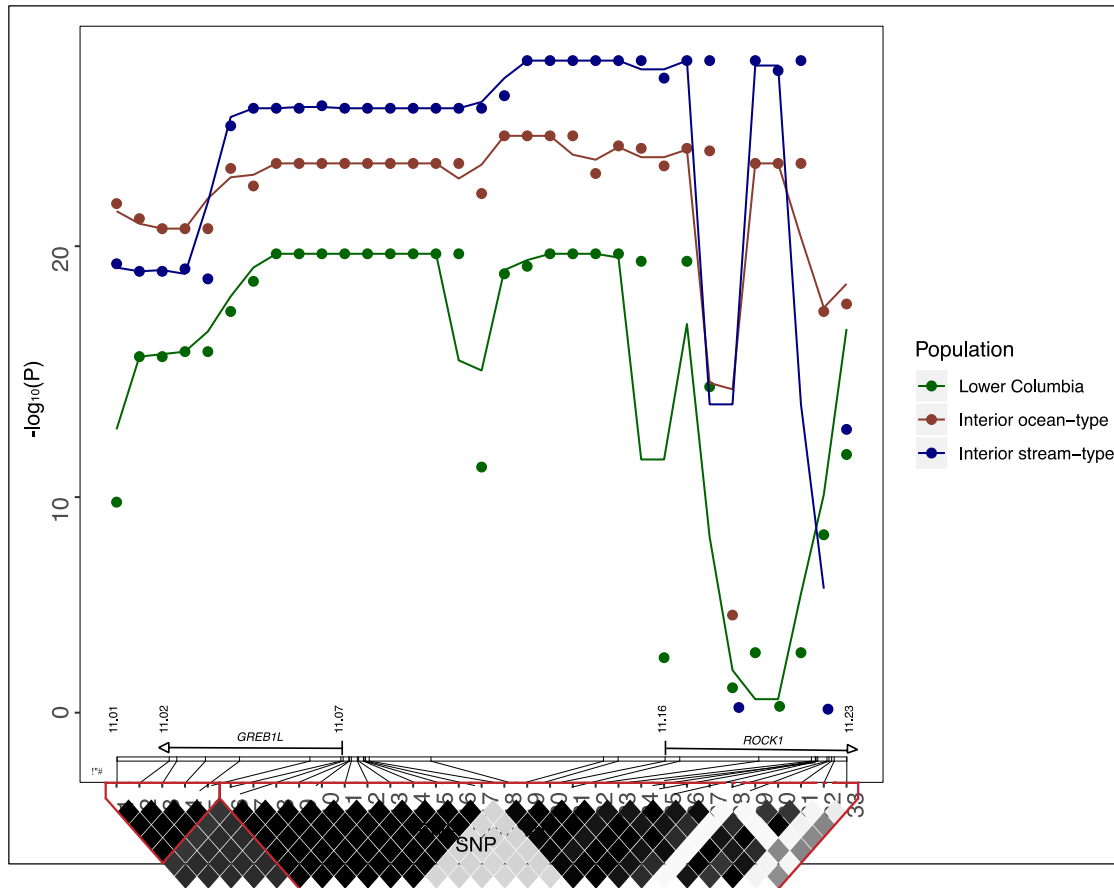


Figure 44. Association of each candidate SNP marker with migration timing within three lineages of Chinook Salmon. The colored lines for each lineage represent sliding windows for consecutive pairs of SNPs along the candidate genomic region. SNP number on the x-axis corresponds to the 33 candidate markers, while the genome position of markers on chromosome 28 (NCBI accession GCA_002831465.1) is depicted in the gene diagrams above the x-axis. Y-axis represents $-\log_{10}(\text{FDR-corrected p-value})$.

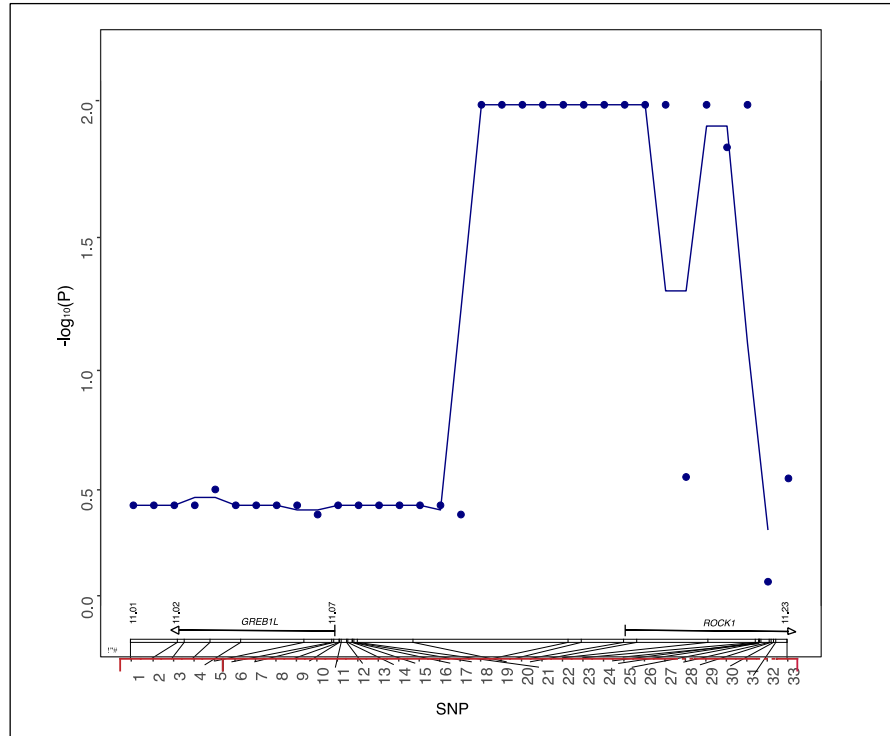


Figure 45. Association of each candidate SNP marker with fitness within the Interior stream-type population. The lines represent sliding windows for consecutive pairs of SNPs along the candidate genomic region. SNP number on the x-axis corresponds to the 33 candidate markers, while the genome position of markers on chromosome 28 (NCBI accession GCA_002831465.1) is depicted in the gene diagrams above the x-axis. Y-axis represents $-\log_{10}(\text{FDR-corrected p-value})$.

Age at maturity in Chinook salmon appears to be a polygenic trait but genes of greatest effect differ between sexes (females = OPN4, males = TMEM19; Figure 46; Micheletti and Narum 2018b). We continue to investigate ocean age/size at maturity for steelhead with whole genome resequencing data for distinct 1-ocean vs 2-ocean phenotypes in multiple populations. Disease resistance in *O. mykiss* appears to be highly complex with several genes involved (Campbell and Narum 2015).

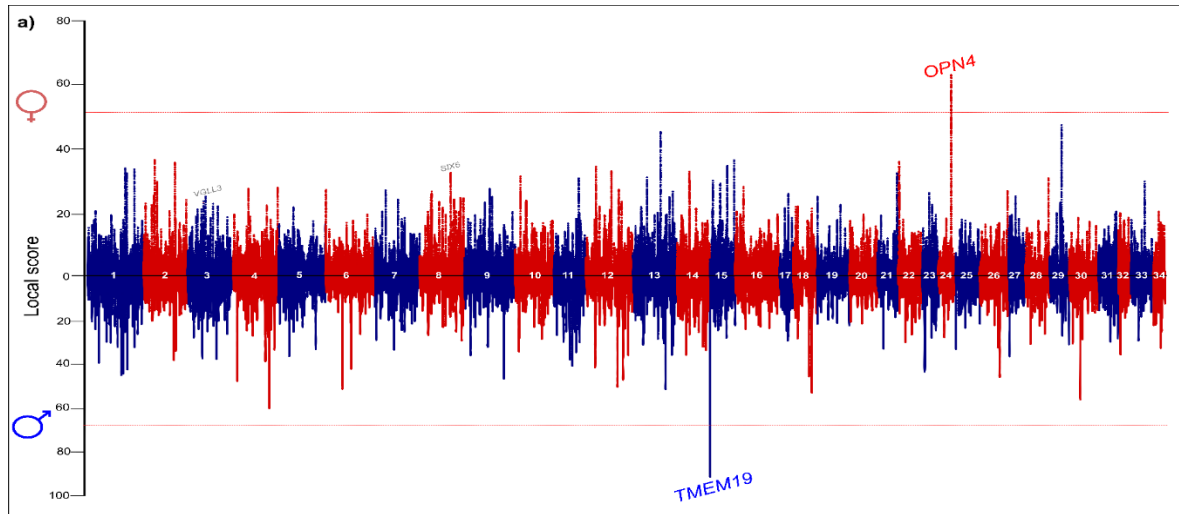


Figure 46. Manhattan plot (from Micheletti and Narum 2018b) illustrating differentiated regions within age classes of females (top; 4- vs. 5-year-olds) and males (bottom; 3- vs. 4- vs. 5-year-olds). Red dashed lines indicate $\alpha = 0.01$ significance threshold for the local score test with Bonferroni correction. Only the opsin 4 (OPN4) gene in females and transmembrane protein 19 (TMEM19) in males were significant using Bonferroni corrected thresholds. Relative locations of the VGLL3 (Ch3) and SIX6 (Chr8) genes which correspond to age at maturity in Atlantic salmon, are displayed in grey and were not significant.

Finally, thermal adaptation has evolved in desert strains of *O. mykiss* (Narum et al. 2010) so that fish have higher thermal tolerance and increased capacity to deliver oxygen to tissues by achieving higher maximum heart rate than montane fish (Chen et al. 2018a; Chen et al. 2018b). The genetic basis for thermal adaptation includes response of heat shock proteins (Narum et al. 2013; Narum et al. 2015; Chen et al. 2018a; Chen et al. 2018b) but also other genes involved in efficient uptake oxygen (Garvin et al. 2015; Narum et al. 2015; Chen et al. 2018a; Chen et al. 2018b). As these candidate genes for traits have begun to be identified (Figure 50; Chen et al. 2018a), SNP markers from these regions are being incorporated in standard genotyping panels with GTseq in order to validate and monitor genetic variation for these traits in large numbers of individuals. Additionally, thermal tolerance has begun to be evaluated in test populations of anadromous steelhead and Chinook salmon following similar measurements of phenotypes as used for redband trout.

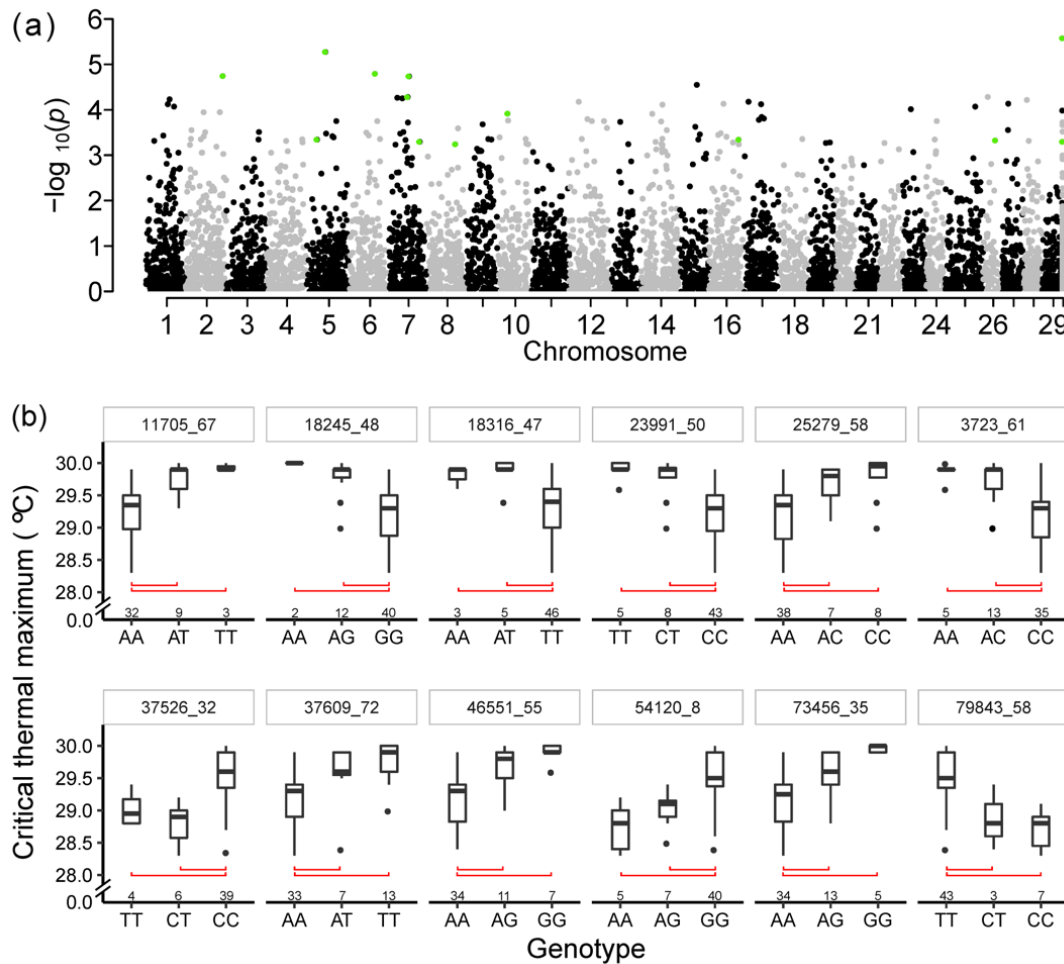


Figure 47. Results from Chen et al. (2018a) that demonstrate association of loci with thermal tolerance in natural populations of redband trout from desert, cool montane, and cold montane environments. Outlier loci and the association with critical thermal maximum (CTMAX). (a) Manhattan plot of calibrated p values, which were derived from the median z-score of results from Lositan, outFLANK and PCAdapt (François et al., 2016). Outliers identified by at least two analyses are in green. (b) significant associations between CTMAX and genotypes of candidate outlier loci (locus name above each panel). Red brackets indicate significant differences at the level of $\alpha=0.05$ in one-way ANOVA on ranks with subsequent Dunn's post-hoc test. Numbers above genotypes represent the sample size. Results published in Chen et al. (2018a).

Summary of accomplishments & lessons learned:

Steady progress has been made to identify adaptive genomic variation and genetic basis for phenotypic traits in steelhead and Chinook salmon as reflected in 17 publications: Narum et al. 2008a; Narum et al. 2010; Narum and Campbell 2010; Narum et al. 2011; Hess and Narum 2011; Hecht et al. 2013; Narum et al. 2013; Campbell and Narum 2015; Garvin et al. 2015; Narum et al. 2015; Hess et al. 2016; Chen et al. 2018a; Chen et al. 2018b; Micheletti and Narum 2018; Micheletti et al. 2018b; Micheletti et al. 2018c; Narum et al. 2018).

Candidate genes for several traits have been identified in both steelhead and Chinook salmon. As these candidate genes for traits have begun to be identified, SNP markers from these regions are being incorporated in standard genotyping panels with GTseq to validate and monitor genetic variation for these traits in large numbers of individuals.

2695

2696 **Synthesis of Findings: Discussion/Conclusions**

2697 ***Fish Population RM&E***

2698

2699 **Management questions/decisions and anticipated outcomes**

2700 *Application of research results:*

2701 Inclusion of candidate markers associated with specific traits allows more detailed
2702 genetic monitoring of stocks in the Columbia Basin. Extensive programs are in place that
2703 enable genetic identification of the origin of individual fish, but candidate markers from
2704 this study also provide the ability to monitor genetic variation for specific traits that are
2705 expected to be necessary to maintain life history variation for long term persistence of
2706 populations.

2707

2708 Water temperatures are predicted to increase in this century, e.g. approximately 0.27°C
2709 per decade for streams where salmonids are distributed. Thus, it is questionable whether
2710 species and populations will be able to adapt to future environmental changes, especially
2711 for freshwater ectotherms with limited migratory opportunities. Local extirpation events
2712 might occur if populations experience extreme temperatures above their maximum
2713 adaptive capacity from existing genomic variation. According to our predictions based on
2714 standing genetic variation at adaptive loci, natural populations appear to have some
2715 capacity to evolve a higher mean CT_{MAX} to meet challenges of warmer conditions (Figure
2716 51). However, populations that currently live in warm environments may have a narrow
2717 safety margin, and therefore are more vulnerable and may need conservation attention.
2718 Thus, phenotypic plasticity and behavioral thermoregulation, such as seeking thermal
2719 refugia (e.g. deep pools, cool springs and upwelling groundwater), will become critically
2720 important for them to temporarily survive or avoid extreme temperatures in the future. To
2721 predict the rate and limits of evolutionary adaptation more acutely in the future, more
2722 advanced niche models need to incorporate additional factors such as the intensity of
2723 selection, effective population size, heritability and phenotypic plasticity. This
2724 information can be incorporated into robust adaptive networks that include a broad
2725 portfolio of adaptive diversity, connectivity, and meta-population scale management for
2726 long-term persistence.

2727

2728 Markers that are associated with specific phenotypic traits will also enable monitoring of
2729 genetic variation for traits that are considered important to managers. In particular,
2730 markers have been developed for premature vs. mature arrival to spawning grounds (e.g.,
2731 migration/maturation timing) to monitor genetic variation for this trait in the Columbia
2732 Basin.

2733

2734

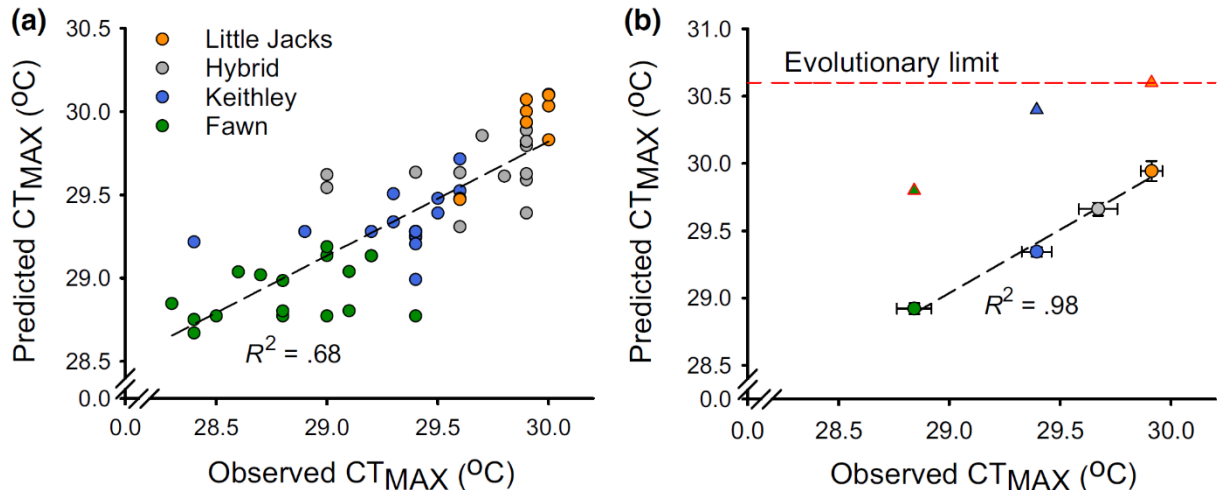


Figure 48. Critical thermal maximum (CTMAX) prediction (from Chen et al. 2018a). (a) Correlation between observed and predicted individual CTMAX values. (b) Prediction of population mean CTMAX (SEM) and evolutionary limits for populations (triangle) and species (dash line).

Project timeline

This project began in 2009 with studies that utilized genetic data available at the time. As genomic methods have advanced over the last decade, more intensive genomic tools have enabled discovery of adaptive genetic variation and the genetic basis for specific phenotypic traits. Work is ongoing to discover adaptive variation and validate the genetic basis for phenotypic variation that is necessary for long-term persistence of salmonids. Thus, the project does not have a clear end date as more work is necessary into the future.

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Section 6: Sturgeon genetics

Introduction

White sturgeon are a long-lived amphidromous species that historically ranged throughout the Columbia River Basin, occurring as broadly distributed, overlapping meta-populations (Parsley 2007; Beamesderfer et al. 2012). Overfishing is believed to have contributed heavily to decreased productivity and declining population abundances as early as the late 1880's (Mallette 2008). Moreover, impacts on white sturgeon populations and trending declines in abundances have been linked to factors associated with the construction of dams (e.g., the hydropower system) in the mainstem Columbia River and its tributaries. Impounding of the Columbia River has significantly fragmented populations (see Nelson et al. 2013). Regional flow regimes and hydrology have been altered and diminished from their natural states (Barton et al. 2010), affecting availability and quality of preferred spawning and rearing habitat for white sturgeon over a large geographic range (Parsley et al. 1993; Rien et al. 2005; Parsley 2007). Most notably, dams have greatly restricted migration through the Columbia River corridor, and fish passage has not provided the same benefit for sturgeon that has been afforded to salmon species (Beamesderfer et al. 2011; Beamesderfer et al. 2012). By some estimates, as many as 24 functionally discrete white sturgeon populations have arisen as a result (Parsley 2007). It is also important to recognize that altered habitat has severely impacted recruitment, predator and prey interactions, and population genetic variability throughout much of the Columbia River Basin (Nelson et al. 2012). Considerable monitoring and evaluation efforts have been initiated in the Columbia (KTOI 2007; Drauch-Schreier et al. 2012; Drauch-Schreier et al. 2013), and Snake River region (IPC 2005), where sturgeon populations have experienced some of the greatest declines, and where little to no recruitment has been observed in recent years. There have been similar and concerted attempts to characterize white sturgeon population status where relatively larger numbers of sturgeon still persist (e.g., Bonneville Reservoir and the lower Columbia River) but where long-term impacts to habitat and productivity remain a concern for long-term viability and genetic diversity (Mallette 2008; Chapman and Jones 2010).

Since 2008, Tribal managers and CRITFC scientists have been monitoring the genetic population structure of white sturgeon residing in reservoirs impounded by Bonneville Dam, The Dalles Dam, John Day Dam and McNary Dam in the Middle Columbia River. A long-term monitoring effort was initiated in order to assess previously described risks to contemporary population status, including concerns for limited migration potential, declining or low genetic diversity, small effective population sizes, and poor productivity. This report provides a summary of ongoing efforts in 2019 to understand of local demographics and resolve population structure. Previously, we reported on our success in overcoming the octoploid nature of the white sturgeon genome, which created difficulties in scoring traditional microsatellite markers, by developing a panel of single nucleotide markers (SNPs) that we are able to genotype as functionally-tetraploid markers. Because the inheritance pattern of these markers is more interpretable than data from microsatellite markers, these data are more amenable to a wider variety of genetic analyses.

Methods

Our SNP panel consists of 325 loci that met or exceeded quality control criteria, including minor allele frequencies above 5%, and at least 80% genotyping success among samples (Matala et al. 2017). The SNPs were designed using extended sequence data, and the panel was genotyped using the GT-seq high throughput sequencing method (Campbell et al. 2015;

<https://www.monitoringresources.org/Document/Method/Details/5446>). Genotypes of white sturgeon segregate into five distinct clusters indicative of four alleles at each locus (tetraploidy) (e.g. AAAA, AAAB, AABB, ABBB, BBBB) (**Figure 49**).

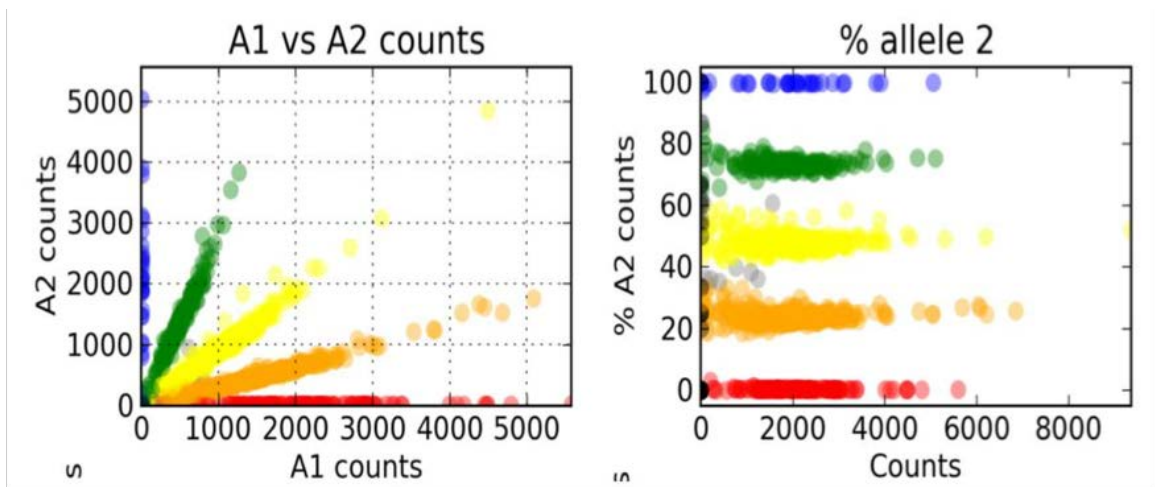


Figure 49. Example of a tetraploid locus GT-seq plot.

This genotyping panel was tested using a set of known parent-offspring pairs from 3 reciprocal pairings (3 males x 3 females) contributed by our collaborator at the University of California-Davis (Dr. Andrea Drauch-Schreier). Using these known relationships, we were able to establish that the majority of loci exhibited the expected mendelian inheritance patterns, while flagging a handful of loci for re-evaluation (**Figure 50**). Though statistical parentage assignment remains challenging (see Discussion), these data also appeared promising for identifying the most likely parents among a set of candidates.

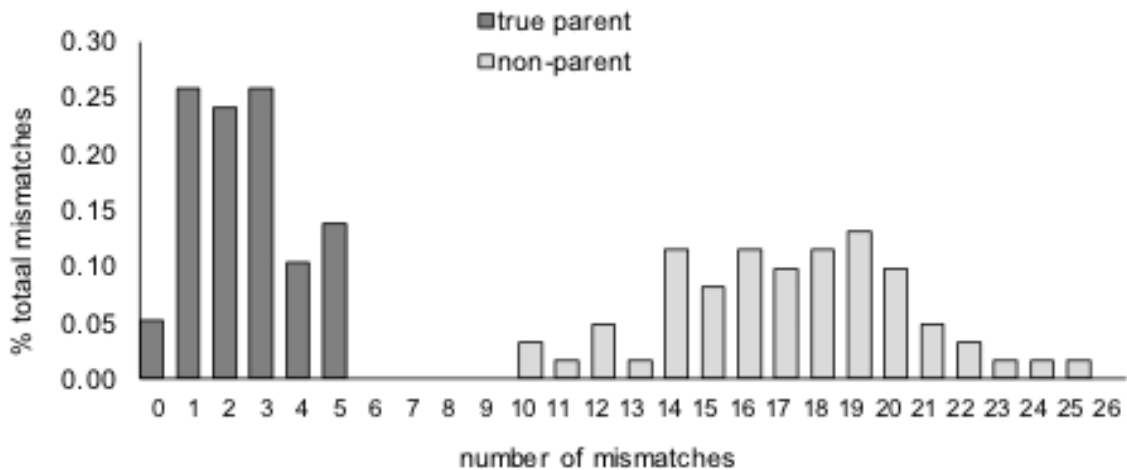


Figure 50. Distribution of proportional discrepancies (numbers of mismatched loci) observed among all offspring assigned to a non-parent pair compared to offspring assigned to a known (i.e. true) parent pair.

Given the success in developing this genotyping panel (described in more detail in our 2018 report), in 2019, in collaboration with sturgeon experts at Cramer Fish Sciences (P. Anders), we

undertook to organize and expand our tissue holdings for white sturgeon. We were specifically interested to identify representation in our holdings sufficient to meet our continuing objectives, including understanding the relationship of genetic diversity to local adaptation and evolutionary potential in the context of regional and local population structure, as well as local demographic trends in recruitment and spawning success, their relationship to recent environmental conditions, and their amenability to stock supplementation.

Results:

Our tissue holdings include 10,697 unique samples from white sturgeon from the Columbia Basin, as well as 116 from other Pacific versants. We categorized these based on the Columbia River reach from which they were collected (**Figure 50**), and where known, the life stage of the individual (Table 68).

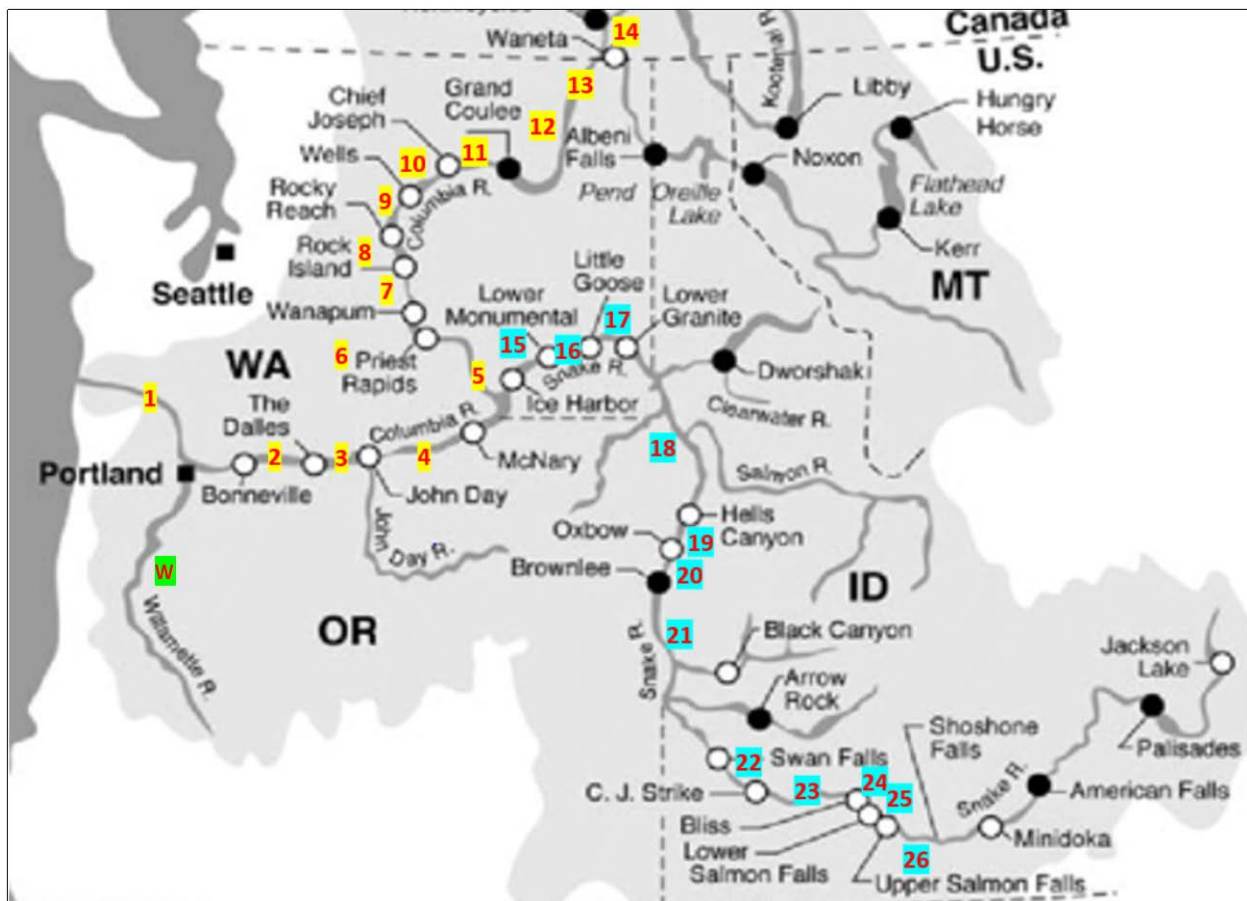


Figure 51. Locations in the Columbia Basin where white sturgeon are or historically were present.

Table 68. White sturgeon from the Columbia Basin in the CRITFC collection. Life stages are defined as Adult (>136cm), Sub-adult (100-136cm), and Juvenile (<100cm).

Loc	Region	Reach	Adult	Sub-adult	Juvenile	Unk	Totals
1	Lower Columbia River	Lower Columba River	159	49	813	41	1,062
2	Lower River Pools	Bonneville Pool	78	6	2,931	342	3,357
3		The Dalles Pool	46	1	1,800	56	1,903
4		John Day Pool	606	492	1,219	851	3,168
5	Mid-Columbia Pools	McNary Pool	50	68	56	53	227
6		Priest Rapids Pool	0	0	0	0	0
7		Wanapum Pool	0	0	0	0	0
8	Upper-Mid C Pools	Rock Island Pool	0	0	0	0	0
9		Rocky Reach Pool	3	6	28	0	37
10		Wells Pool	0	0	0	0	0
11		Chief Joseph Pool	0	0	0	0	0
12	Upper Columbia	Grand Coulee Reservoir	99	1	126	0	226
13		Upper Columbia River	0	0	0	0	0
14		Upper Columbia (BC)	0	0	0	0	0
15	Lower Snake	Ice Harbor Pool	25	33	65	4	127
16		Lower Monumental Pool	67	52	25	0	144
17		Little Goose Pool	62	42	27	0	131
18		Lower Granite Pool	24	10	32	30	96
19	Middle Snake	Hells Canyon Reservoir	0	0	0	0	0
20		Oxbow Reservoir	0	0	0	0	0
21		Brownlee Reservoir	0	0	0	0	0
22		Swan Falls Reservoir	0	0	0	0	0
23		CJ Strike Reservoir	11	4	0	78	93
24	Upper Snake	Bliss Reservoir	0	0	0	0	0
25		Lower Salmon Falls Res.	0	0	0	0	0
26		Upper Salmon Falls Res.	0	0	0	112	112
W	Willamette	Santiam River	0	0	0	14	14
			1,230	764	7,122	1,581	10,697

Tallying using meta-data characteristics in this manner allowed us to assess the strengths of this collection, as well as to identify areas for additional sampling and tissue loans, although in general the abundance of samples in our collection reflects where there remain moderate to robust populations of white sturgeon in the Columbia Basin (Beamesderfer, et al. 2012). From this parsing, we were able to identify a subset of 3,006 samples to genotype in the 2020 calendar year that will allow us to assess, using the more robust 325 marker panel, population genetic structure in the Columbia Basin, and the Columbia relative to other drainages, as well as variations in genetic diversity across life stages indicative of differing and potentially problematic patterns of recruitment (Table 69).

Table 69. White sturgeon targeted for genotyping in calendar year 2020. Life stages are defined as above, except that fish less than 30cm are identified as young-of-year (YOY).

	Region	Reach	Adult	Sub-Adults + Juveniles	YOY	Unk	Totals
1	Below Bonneville	Lower Columbia River	85	68	223	3	379
2	Lower Columbia	Bonneville	36	5	264	0	305
3	Lower Columbia	The Dalles	18	8	327	1	354
4	Lower Columbia	John Day	249	421	81	142	893
5	Middle Columbia	McNary	46	97	14	55	212
9	Upper-Mid Columbia	Rocky Reach	3	34	0	0	37
12	Upper Columbia	Grand Coulee/FDR	99	27	100	0	226
15	lower Snake	Ice Harbor	26	87	0	4	117
16	lower Snake	Little Goose	66	54	0	0	120
17	lower Snake	Lower Monumental	68	74	0	0	142
18	lower Snake	Lower Granite	22	3	0	18	43
23	middle Snake	CJ Strike	8	1	0	75	84
26	upper Snake	Upper Salmon Falls	0	0	0	96	96
W	Willamette	Santiam	0	0	0	14	14
	Outside Columbia	Nechako River	0	0	0	5	5
	Outside Columbia	Sacramento-San Joaquin River	6	0	0	3	9
	Outside Columbia	Fraser River	0	0	0	10	10

Discussion:

Our research is guided by previous results which indicate that white sturgeon in the Middle Columbia River have experienced a significant genetic bottleneck and decreased diversity as a result of dramatically restricted migration, altered flow regimes, and loss of spawning and rearing habitat. In addition, little to no genetic differentiation between impoundments in the Middle Columbia River has been observed, although a regional distinction between Snake River and Columbia River populations is evident (Matala et al. 2017). The accumulation of genetic monitoring results (2008-2018) for White Surgeon in the Middle Columbia River and Snake River are similar to evaluations of population structure reported for similar large systems like the Fraser River in British Columbia Canada. Current demographic trends and genetic variation may be a consequence of diminished rearing and/or spawning habitat coincident with stream fragmentation and limited connectivity. Ongoing efforts for this project are designed to further resolve the dynamics of recruitment and population demographic trends in the context of regional population structure and local adaptation and evolutionary potential, particularly in the John Day and Grand Coulee Reservoirs, where Tribal efforts are underway to identify efficient stock supplementation techniques (Table 69).

To this end, a robust parentage baseline will provide confident estimates of the size of spawning populations (i.e. numbers of adults contributing to recruitment) based on levels of kinship among juvenile age groups. However, one of the ongoing challenges with this objective is the nature of genetic data from white sturgeon. Although technically octoploid (8 of each chromosome), following an evolutionarily recent genome duplication from an putatively tetraploid ancestral state (4 copies of each chromosome), the white sturgeon appears to have regained a functional tetraploid status, as indicated by the mendelian segregation of four alleles at the majority of SNP loci (**Figure 49**). Nonetheless, while the optimization of markers accurately rendering genotypes

of the functional tetraploid chromosomal segments empowers our ability to glean information about population genetic structure and diversity, the majority of statistical parentage programs remain designed to process only diploid data (two copies of each chromosome, like humans). While we have garnered some success in identifying likely parent candidates using ad hoc (e.g. mismatch distribution) methods (**Figure 50**), we continue to develop and explore more robust methods to fully utilize the information content of tetraploid genotypes. Robust information on recruitment dynamics will be valuable to managers concerned with reversing decreasing trends in productivity, and in hatchery operations focused on maximizing genetic variation and family contributions in supplementation or translocation programs. Distance and numbers of barriers that currently sequester populations in the Snake River region and Middle Columbia may profoundly influence genetic variability. Some of the highest levels of genetic diversity in our long-term analyses were observed in the Lower Columbia River population of relatively high abundance. However, where historical migration between regions has been restricted, the effective isolation of larger and older mature fish means the opportunities for populations to repopulate or recolonize will take decades following recruitment. Therefore, understanding the origins and abundances of local spawners is vital for estimating demographic trends, the potential to improve genetic variability, and the efficacy of supplementation. Targeting of both YOY and older (mature) fish will continue in subsequent years to allow for implementation of additional kinds of proposed analyses defined in the overall objectives of this project. In addition, our continuing efforts with white sturgeon include the development of a draft genome assembly. While extensive whole genome data has provided draft contig sequences, the majority of the genome remains insufficiently anchored for publication due to the challenges of scaffolding a high repetitive (octoploid) genome. To overcome this, we are working closely with collaborators to generate a linkage map from a reduced ploidy line of white sturgeon, which will assist to order or orient (anchor) smaller scaffolds into chromosomes. The draft genome, once complete, will allow us to use genome-level sequencing data to survey genetic variants underlying phenotypic traits and local adaptation, a type of analysis that the lab has had extensive success completing for other Columbia Basin species (e.g. Micheletti et al. 2018). Since White Sturgeon are long lived and require many years to reach sexual maturity, the value of data generated by this project are largely defined by the project's long-term implementation. It began in 2008 and continues to present, with samples collected and archived on an annual basis. Among other things, this affords the opportunity to monitor cohorts through various age-classes or life history stages. It also provides more robust analysis as data accumulates. For example, fish that were sampled as subadults in 2008 are reaching (or have reached) sexual maturity and are available to screen as potential parents of juvenile fish sampled in recent or upcoming years. Work on our most current objective (genome sequencing to discover a sex-linked marker for gender ID) has only just begun, and we anticipate long-term applicability of the data produced, particularly with continued collaborations and data sharing with co-managers in the basin.

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Overall Conclusion

This project combines four inter-related studies from the Fish & Wildlife Program Accords that address the following current and future objectives: 1) discover and evaluate SNP markers in salmon and steelhead and other anadromous fishes; 2) expand and create genetic baselines for multiple species including Chinook salmon, steelhead, sockeye salmon and kokanee, and coho salmon; 3) implement Genetic Stock Identification (GSI)/PBT programs for mainstem Chinook salmon, sockeye salmon, and steelhead fisheries and 4) GSI/PBT of fish passing Bonneville Dam (steelhead, sockeye, and Chinook salmon).

As described in Section 1, SNP panels continue to be expanded with GTseq that enables genotyping large sample sizes (>157,000 fish genotyped in 2019). This genotyping protocol has greatly increased our laboratory's efficiency by allowing large numbers of fish to be genotyped with large numbers of SNP loci but at lower costs. For genetic baseline expansion (Objective 2), PBT hatcheries above Bonneville were genotyped to enable more thorough assignment of hatchery origin fish. In addition, GSI baselines are being developed with whole genome resequencing to provide allele frequency estimates for millions of SNPs in Chinook salmon and steelhead. SNPs identified through current efforts involving whole genome resequencing will be useful in characterizing genetic diversity of hatchery and wild Chinook salmon and steelhead stocks. This study included two broad applications of stock identification; namely, stock composition of fisheries for Chinook salmon, sockeye salmon, and steelhead (Objective 3), and stock composition of Chinook salmon, sockeye salmon, and steelhead passing Bonneville Dam (Objective 4). Chinook salmon and steelhead fishery applications of GSI were integrated with the new genetic technology of parentage based tagging (PBT). The challenge imposed by long histories of exogenous stock transfers from specific hatchery programs often prevents effective application of GSI in assigning hatchery fish. However, as the role of PBT is expanding to tag all hatchery fish, the role of GSI will be focused on identifying stocks of natural-origin fish.

Our GSI analyses of harvest included stock composition results for the spring, summer, and fall management periods of Chinook salmon fisheries in the lower Columbia River mainstem. Sockeye salmon fisheries were analyzed and our stock composition results will provide additional information to managers of these fisheries. However, the sockeye salmon results indicate an increase in sample size may be warranted to make accurate estimates of rare stocks such as Snake River sockeye salmon.

For Objective 4, we used a combination of GSI and PBT to estimate run-timing distributions and relative abundance of hatchery and wild Chinook salmon and steelhead stocks in 2019. For sockeye salmon, we used GSI to estimate relative stock abundance and run-timing distributions. The stock-specific data on abundance and run-timing of these species were used as a context for interpreting harvest stock composition. We continue timely post-season reporting of genetic analysis of Chinook, steelhead and Sockeye Salmon at Bonneville Dam in 2019, but we also continue to provide in-season results at bi-weekly intervals during the spring and summer management periods of Chinook Salmon, and approximately monthly for the summer and fall runs of steelhead and Chinook Salmon. The timely updates of the genetic analyses for these runs and species of Columbia River fishes improves their utility for fisheries managers such as the Technical Advisory Committee (TAC).

For Objective 5 on local adaptation, candidate genes for several traits have been identified in both steelhead and Chinook salmon. As these candidate genes for traits have begun

to be identified, SNP markers from these regions are being incorporated in standard genotyping panels with GTseq to validate and monitor genetic variation for these traits in large numbers of individuals. Inclusion of candidate markers associated with specific traits allows more detailed genetic monitoring of stocks in the Columbia Basin. Extensive programs are in place that enable genetic identification of the origin of individual fish, but candidate markers from this study also provide the ability to monitor genetic variation for specific traits that are expected to be necessary to maintain life history variation for long term persistence of populations.

For Objective 6 on white sturgeon genetics, the accumulation of genetic monitoring results (2008-2018) for White Surgeon in the Middle Columbia River and Snake River are similar to evaluations of population structure reported for similar large systems like the Fraser River in British Columbia Canada. Current demographic trends and genetic variation may be a consequence of diminished rearing and/or spawning habitat coincident with stream fragmentation and limited connectivity. Ongoing efforts for this project are designed to utilize SNP markers that were developed for this species to further resolve regional population structure and parentage analyses to support tribal efforts to supplement white sturgeon in the Columbia Basin.