

Columbia River Project Water Use Plan

Columbia White Sturgeon Management Plan

White Sturgeon Conservation Aquaculture – 2022 Annual Report

CLBWORKS-25: Mid-Columbia River White Sturgeon Conservation Aquaculture

CLBWORKS-34: Lower Columbia River White Sturgeon Conservation Aquaculture

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BC Hydro and Power Authority

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Mid and Lower Columbia River White Sturgeon Conservation Aquaculture Report: 2022

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Society of BC**

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1. Background

The population of White Sturgeon in the Canadian portion of the Columbia River has been undergoing recruitment failure for several decades (Hildebrand and Parsley 2014). This was recognized as a critical issue for this population in the early 1990's and resulted in the establishment of the Upper Columbia White Sturgeon Recovery Initiative (UCWSRI; <http://uppercolumbiasturgeon.org>) in 2000 and the population being listed as endangered under the Species at Risk Act (SARA) in 2006. Original estimates, based on annual levels of natural recruitment being insufficient for maintaining a self-sustaining population, suggested that the population would become functionally extinct by 2044 (UCWSRI 2002; Hildebrand and Parsley 2014). Accordingly, intervention and monitoring were deemed essential to preclude extinction. An integral part of the original recovery plan was the initiation of a conservation aquaculture program designed to support the population until such time as stock abundance/age structure and habitat conditions can support a self-sustaining population.

The white sturgeon conservation aquaculture program was initiated in 2001 in the lower Columbia River Canada, encompassing 57.0 km of the Columbia River from Hugh L. Keenleyside (HLK) Dam near Castlegar British Columbia to the international border. A parallel program was initiated in the US in 2003, to produce juvenile sturgeon for release into Lake Roosevelt. Finally, in 2007, the program was extended to the Middle Columbia River Canada, which is a 235 km section of the Columbia from HLK to Revelstoke Dam and includes the Arrow Lakes Reservoir. Annual releases of sub-yearlings have occurred into all locations since their inception. The key overarching objectives of the aquaculture program were originally to prevent extirpation and retain the genetic diversity of the wild population. In order to achieve these objectives, the program followed an adaptive management approach that continually incorporated new methods and key results into the decision-making process. The adaptive management of the program was conducted in partnership with UCWSRI partners from both Canada and the US that included agencies, First Nations, industry, and other stakeholders. A major transition in

the program occurred in 2014, when the use of adult broodstock as an egg source was replaced with capture of wild eggs and larvae in order to maximize genetic diversity.

The main objective of this report is to provide a summary of the Canadian conservation aquaculture activities for the 2021 brood year. Further details about the longer-term methods and results can be found within annual and comprehensive reports (details in FFSBC 2008-2020 available at

https://www.bchydro.com/toolbar/about/sustainability/conservation/water_use_planning/southern_in_terior/columbia_river/columbia-sturgeon.html).

2. Methods

Background of Wild Origin Approach

The conservation sturgeon aquaculture program successfully reared and released 164,423 sturgeon originating from captured broodstock between 2002 and 2015 (brood years 2001 to 2014). However, survival of hatchery-origin juveniles in the upper Columbia River population has been higher than originally predicted, with more than 20,000 individuals estimated to be at large in the population (BC Hydro 2018; Crossman et al. 2023). Within the hatchery population, certain year classes that benefited from high survival due to large release size are now estimated to be in higher abundance than the existing wild population (~3,000 mature individuals). Of further uncertainty was whether within year class genetic diversity has been reduced relative to the expected diversity released from the hatchery due to disproportionate survival among maternal family groups post stocking. To ensure methods for progeny collection were as genetically robust as possible, a large-scale adaptive management change was made to cease use of broodstock and rely completely on capture of wild progeny. This had been developed for lake sturgeon conservation aquaculture (Crossman et al. 2011) to improve genetic diversity, and work in the Columbia demonstrated it was feasible to capture large

numbers of wild larvae and eggs downstream of known spawning locations (primarily Waneta). In the US, the conservation aquaculture program adopted a wild-origin approach in 2011, where feeding larvae migrating downstream from spawning locations were collected and transferred to the hatchery for rearing. In Canada, work by Jay et al. (2014) demonstrated an improvement in genetic diversity and, given the objectives of the overall program to rebuild abundance and maintain genetic diversity, the use of wild progeny was deemed as the optimal source for conservation aquaculture program going forward. Therefore, in 2014 a mobile streamside rearing facility was piloted as an incubation method for wild eggs in conjunction with the traditional broodstock capture. Following 2014 the broodstock program was discontinued and the wild-origin approach has continued to date.

Study Area

The Canadian portion of the Columbia River is described spatially at several different levels. Most broadly, the river is divided into two sections above and below Hugh Keenleyside (HLK) dam which controls the water level of Arrow Lakes. Directly adjacent on river left to the HLK facility is the Arrow Lakes Hydro generating station (ALH). Sturgeon populations above and below HLK are treated as separate populations and have independent recovery and aquaculture release targets (see below). Specific sites within the Canadian portion of the Columbia River are further described by a linear river kilometer relative to HLK dam. The 56km of river below HLK is also categorized into 5 equal zones (11.2 km each). Lastly, there are numerous site descriptions and nicknames for specific capture and release locations. The capture and release site names and zones are all shown in Figure 1.

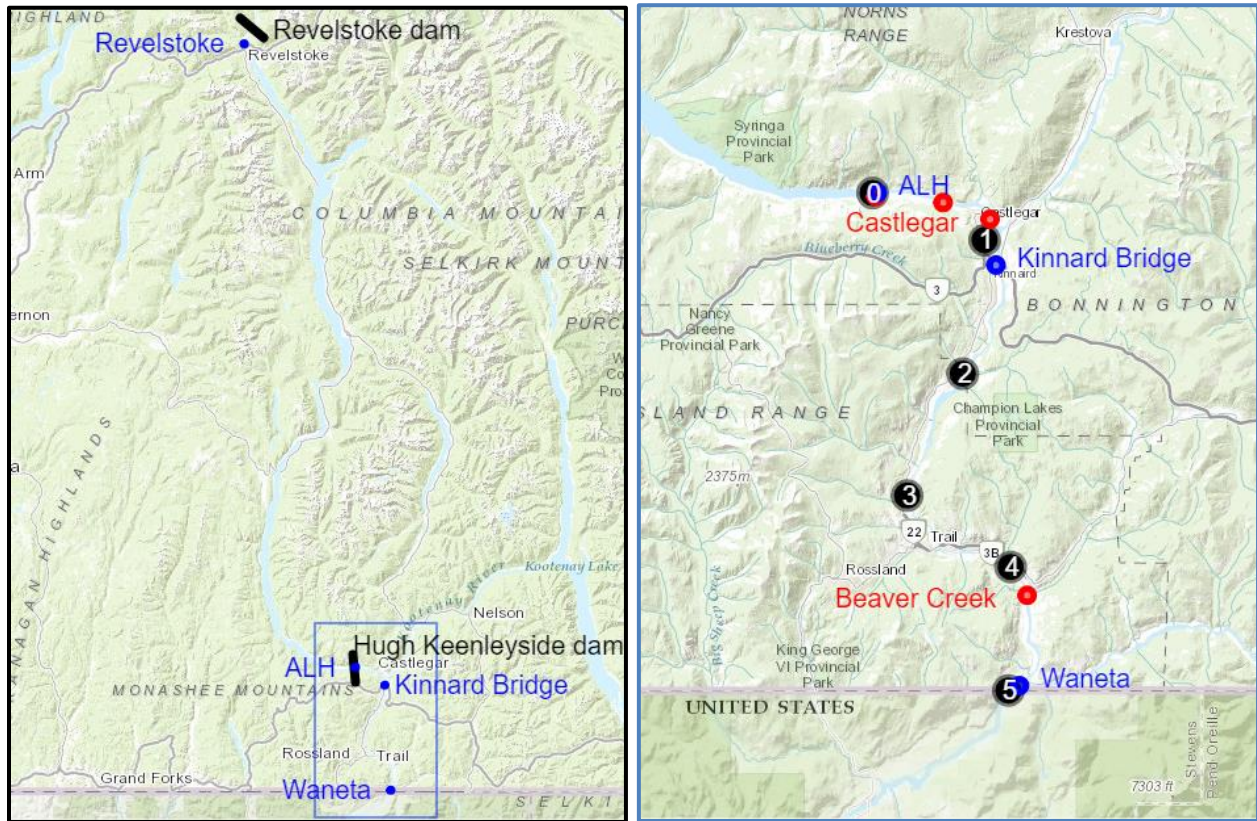


Figure 1. Left panel: Map of the extent of lower Columbia River in Canada and all known White Sturgeon spawning locations where collection of wild-origin embryos or larvae occurs (blue). Blue rectangle refers to the spatial extent of the right panel. Right panel: Map of the spawning sites (blue), juvenile release sites (red), and 5 river zones (black, numbered at downstream limit) for the Columbia River below HLK dam. Note, that fish released to “Castlegar” are spread among three proximal sites (ALH, Robson boat ramp and Millennium Park).

2.1 Early life stage methods

Capture of wild origin sturgeon

Sturgeon are broadcast spawners which allows for early life stages to be sampled with passive sampling gear (Egg mats and Drift nets). The captured contents of egg mats were obtained by carefully inspecting the matt surface and picking off embryos and larvae. Drift net mesh and capsule contents were rinsed into a white bucket, and then transferred in small, diluted

aliquots to several white plastic inspection trays. The trays allowed for contrast and dispersion of debris when searching for the tiny (often sediment covered) embryos. Methods and results of monitoring spawning activity are described in BC Hydro reports (BC Hydro 2023).

Spawning locations

Monitoring of spawning activity occurs at four locations in the Canadian portion of the Columbia River including at Waneta, Kinnaird near Castlegar, downstream of Arrow Lakes Hydro (ALH) and Hugh L. Keenleyside (HLK) Dams, and near Revelstoke (Figure 1). Progeny collected from all spawning locations are brought to the hatchery for rearing. For the 3 sites in the lower Columbia (Waneta, Kinnaird, ALH/HLK), embryos and larvae are incubated in a streamside trailer (details in the following section) prior to transfer to the hatchery. In the 2022 brood year and previous years, progeny collected near Revelstoke were transferred directly to the hatchery from the river as instream or streamside incubation was not possible.

Progeny are grouped by their spawning site of origin and by the spawning event they were collected from. This is done by developmentally staging embryos and larvae in the field at the time of capture and then back calculating their estimated spawning date based on known relationships between development and stream temperatures (Jay et al. 2020). Once at the hatchery, some initial collection groups are combined into larger rearing groups due to space limitations (The maximum number of independent rearing areas is 25, for all brood years present). These groups are only combined if they are groups from the same collection area, and if the spawn event dates are close to each other. Spawn events and therefore their groups would not be combined if they are too far apart. All groups from individual areas are kept separate through the rearing process until they are tagged and therefore become individuals, only then are they combined with other groups. We are able to keep up to 25 groups separate through to the PIT tagging process. If there are more groups than that, or if some of the groups only have a small number of individuals (under ten), then we look at combining like groups. Only small numbers of juvenile sturgeon are required for release, and only a maximum of 500 eggs/larvae are kept from each spawning event, with excess progeny returned to the river.

Streamside incubation trailer

A streamside rearing trailer was created from a 14' by 8' cargo trailer to allow eggs and embryos to incubate at natural stream temperatures. The cargo trailer was set up with an electrical and plumbing system to enable pumping of river water through a series of MacDonald (upweller) jars, that spilled into four troughs with dividers to capture and segregate progeny from separate spawning events (Figure 2).



Figure 2. Streamside Incubation Facility (outfitted cargo trailer) used for rearing wild origin White Sturgeon progeny on the Columbia River. Image from James Crossman

Captured live embryos and larvae were immediately transferred to the streamside incubation facility. Embryos were placed into hatching jars whereas larvae were placed directly into the trough. In both cases they would be segregated based on their estimated spawning event group created from developmental stage and capture location.

Twelve incubation jars (MacDonald Type; J30, Dynamic Aqua-Supply Ltd., Surrey, BC) were available to group embryos by their unique spawning event and location. Water flows are pumped from the Columbia River and maintained at approximately 5 L/min per jar to ensure adequate embryo separation and oxygenation. Upon hatch, yolk-sac larvae were flushed from the hatching jars directly into a segregated section of the rearing trough associated with each

hatching jar (larvae would be placed directly into a section of the trough for their spawning event group). The troughs were supplied with artificial substrate (1" diameter sinking Bio-Spheres; Dynamic Aqua-Supply Ltd. Surrey, BC) which allowed yolk-sac larvae to burrow into interstitial spaces mimicking behavior documented in the wild (McAdam 2011). To reduce sediment in the incubation jars and tanks, water was filtered (254 micron; Spin-Down Separator, Denton, TX) and tanks were cleaned almost daily by purging to remove sediment and waste. All yolk-sac larvae were transported to the Kootenay Sturgeon Hatchery within 7 days of hatch in bags of ambient river water filled with oxygen. Temperature loggers inside the facility recorded air, river water, and facility tank water temperatures.

Transport of Larvae

Larvae need to be in the hatchery for rearing before 10 days post hatch to ensure they can be successfully transitioned to first feeding. FFSBC staff made regular trips to the facility to retrieve larvae and transport them back to KTH to meet the 10 day deadline.

The larvae transport process was as follows: (1) Place two plastic bags (heavy duty clear) in a 10L pail and fill with 4-6 liters of ambient river water; (2) Carefully count and move a single spawn event group of larvae from their trough compartment into the bag; (3) Expel residual air from inner bag, then insert tygon hose from oxygen tank to fill the inner bag with oxygen, then twist and seal with electrical tape, then pull up the outer bag, twist and seal as the inner bag; (4) Places the larvae bag inside a chest cooler containing at least four bubble wrap covered ice packs, close the cooler lid, and place in the truck; (5) Transport to Kootenay hatchery with extra bags and oxygen on hand; (6) Once arriving at the hatchery, float each larvae bag within the destined container (specific to that group) to allow the water temperature in the bag to slowly equilibrate with the container water, then finally cut open the bag and release the fish into their container. As in the streamside incubation facility, each spawning group was placed into a separate container.

2.2 Juvenile rearing

One of the most important aspects of the juvenile rearing process is that each spawn event group (defined by unique location and/or date) must remain segregated throughout the entire rearing process at least until fish can be individually marked (below). Capture of progeny is not equal among spawning events, and some groups are very small (can even be 1 individual) but must still use a separate tank. This is to ensure that at the time of release numbers of progeny are balanced proportionally across all spawning events to the extent possible, maximizing the potential genetic diversity represented.

Larval transition to feeding

The sturgeon larvae remain largely sedentary over the first 10-15 days post hatch while they consume their yolk reserves and continue to develop body parts and organs. The transition to first feeding is a critical life stage for many fish species in both aquaculture and natural settings that can largely determine the relative success of a cohort. White Sturgeon require special attention at this stage.

The troughed larvae start to change behaviour at about day 18 (260 ATUs). Food then can be introduced soon after this behaviour is observed and typically occurs on day 20 (290 ATU). The initial feeding was done by hand with the following process: drain level of pond/trough slightly to expose a damp side, place food onto the wet area on side of trough, allow water level to rise back to previous level so that fish can feed off sides of the trough. In addition to the hand feedings a 24-hour belt feeder was operated to continually dispense a target amount of food as close to the edge of the circular as possible. Each time the pond was drained down for hand feeding (placing feed on the pond walls), the old feed was gently swept out of the pond drain. The type of food used for Sturgeon rearing has varied over the years for a variety of reasons, however, the current mix is presented in Table 1.

Table 1. Diet mixes used to feed different stages of Sturgeon in Kootenay Trout Hatchery from 2013 to present.

Fish weight (g)			Biodiet (pellet size mm)	Krill	Golden Pear
0.2	to	1.0	27% (0.3)	53%	20%
1.0	to	1.7	30% (0.3), 20% (0.5)	40%	10%
1.7	to	6.6	20% (0.3), 70% (0.5)	10%	0%
6.7	to	20.0	40% (0.5), 40% (0.7), 20% (1.0)	0	0
20.0	to	50.0	19% (0.5), 38% (0.7), 38% (1.0), 5% (1.2)	0	0
50.0	to	greater	40% (0.7), 40% (1.0), 20% (1.2)	0	0

Feeding rates were to excess (50% of body weight per day) for first two weeks, and continuous cleaning of the trough was necessary. After the first two weeks, the feed amount was based on temperature and body weight as per Table 2, and the feed sizes also increased to match fish size (starting at 0.3mm and up to 1.2 mm by about day 45). The relationship between body weight and feed rate in Table 6 is a simple power function with a slope of -0.355, and a temperature dependent intercept that corresponds to the row for 1g mean body weight. The early larvae were reared in troughs for about 45 days, and then moved to self-cleaning circular tanks.

Table 2. White Sturgeon feeding rate guidelines as a function of average size and ambient temperature (units are percent of body mass per day).

Mean Body Weight (g)	Temperature (°C)				
	9	11	13	15	17
0.05	21.6	2.7	5.9	4.6	5.2
0.10	16.9	2.9	5.7	4.7	5.2
0.15	14.6	3.1	5.6	4.7	5.1
0.20	13.2	3.2	5.5	4.7	5.1
0.25	12.2	3.3	5.4	4.7	5.1
0.50	9.5	3.6	5.3	4.8	5.1
0.75	8.2	3.8	5.2	4.8	5.1
1.0	7.4	3.9	5.1	4.8	5.1
2.5	5.4	4.4	4.9	4.9	5.1
5.0	4.2	4.8	4.8	5.0	5.0
7.5	3.6	5.0	4.7	5.0	5.0
10.0	3.3	5.2	4.6	5.0	5.0
25.0	2.4	5.9	4.4	5.1	5.0
50.0	1.9	6.4	4.3	5.1	5.0
75.0	1.6	6.7	4.2	5.2	5.0
100.0	1.5	7.0	4.2	5.2	5.0
250.0	1.0	7.8	4.0	5.3	4.9
500	0.8	8.5	3.9	5.3	4.9

Sturgeon larvae develop and grow at rapid rates, which makes first feeding a concentrated and sensitive time period. The fish were minimally handled at this stage, and data on growth and survival rates were not compiled. However, studies presented in a more comprehensive report, BC Hydro 2019 (CLBWORKS 24, 25, 34), show sturgeon exhibit an exponential gain in weight over the first 50 days and a non-linear mortality function. Mortality patterns show substantial

mortality occurs in in the first 21 days (highest mortality rate was between 11 and 21 days), and then stabilizes at a much lower rate after that point.

Juvenile rearing

Once mortality rates appear to have stabilized and increasing tank biomass begins to reach capacity (individuals were approximately 0.2g in size), the fish will be graded by size in order to improve overall growth and survival. The grading of individuals by size into separate tanks reduces overall population densities and ensures smaller individuals can compete for a fair ration of available food. This allows the smaller fish to achieve increased growth rates, so that they catch up to the larger individuals, and the overall group can more uniformly achieve the target release size. The size targets for 2021 brood year production were 200g for fish to be released below Arrow Lakes reservoir, and 300g for releases into Arrow Lakes (Targets section below).

Regardless of size gradation, unique spawning groups were reared separately until fish could be individually marked (see marking and releases below). Juveniles were sampled for size and biomass, as well as assessed individually for disease or deformities. Data is recorded in a group data spreadsheet for all stages prior to pit tagging, and then in an individual data spreadsheet once pit tags (see below) have been applied.

2.3 Juvenile sampling, marking and polyploid testing

The conservation and research basis of the sturgeon conservation aquaculture program requires that the fish be uniquely identifiable when recaptured in the wild. Therefore, prior to release (once the fish reached a size of approximately 50g) each juvenile sturgeon underwent two marking procedures: (1) Each individual was implanted with a PIT tag into the dorsal musculature at the midpoint between the dorsal and lateral scute line inferior to the anterior margin of the dorsal fin. Upon recapture in the wild, PIT tags allow individual level tracking back

to the release records which include data for: brood year, group (defined by egg/larvae capture location and date), release date, release location, release length and weight. (2) All hatchery fish also have a set number/location of scutes removed so that they are quickly visually identifiable in the field. The side of the body from which scutes were removed was changed to match the source/stage that were first reared in the hatchery from broodstock spawn events (left side) to wild origin larvae (right side). All mark locations are depicted in Figure 3. DNA and blood were also taken during the marking process (see Polyploidy testing section below).

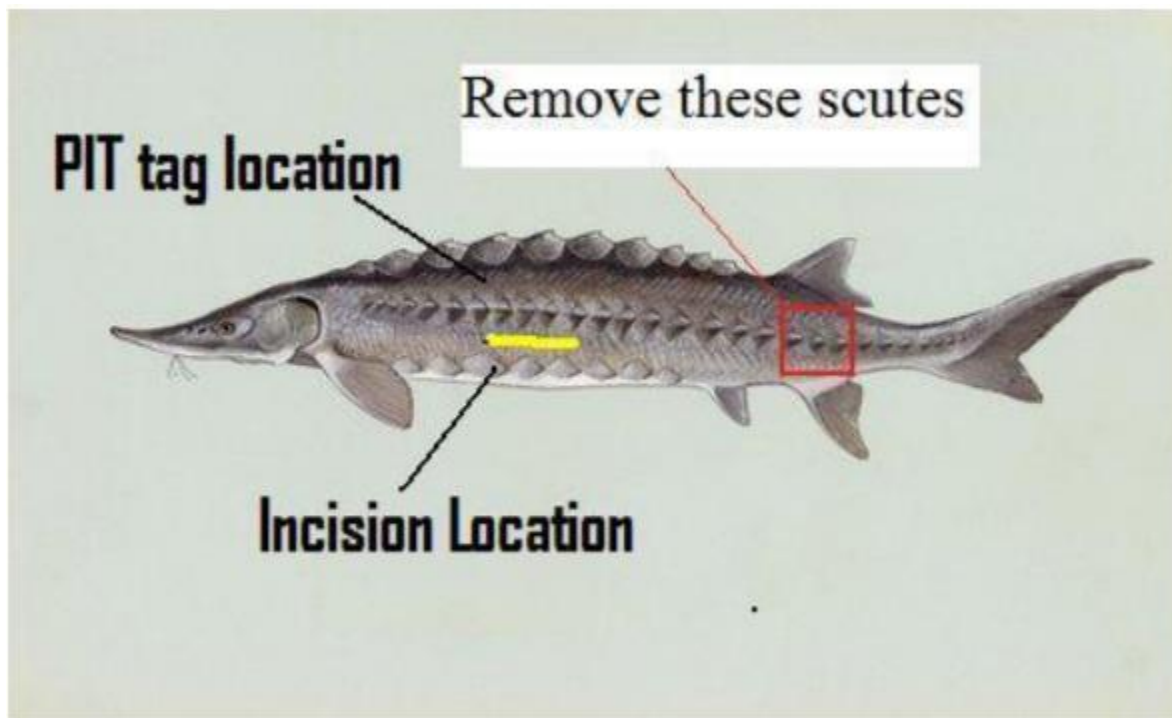


Figure 3. Diagram of physical locations of marks applied to hatchery reared juvenile sturgeon. Incision location refers to the location of acoustic tag insertion in years where acoustic tags were released (image extracted from BC Hydro CLB WORKS-34 2013 report).

Immediately prior to release (within 1 week) all fish were measured for length and weight in order to accurately reflect their size-at-release. A sub-sample of fish were run through an additional disease screening. The juveniles were transported to the release site using FFSBC

transport vehicles according to UCWSRI TWG transport protocols. Sturgeon were strategically released at locations as directed by the technical working group (See releases).

Autopolyploidy

The ploidy (number of sets of chromosomes) of White Sturgeon is typically 8N in Columbia and Kootenay River populations (Schreier et al. 2011, Schreier et al. 2013, Gille et al. 2015; Schreier et al. 2021). However, 12N White Sturgeon were found to occur (at a rate of up to 50%) at the Kootenay Tribe of Idaho's (KTOI) Sturgeon Hatchery in family groups that were derived from their wild broodstock program. The mechanism of this ploidy shift is unknown, but it is hypothesized to occur at the fertilization stage. Given the findings in the Kootenay River White Sturgeon, the Columbia River Technical Working Group directed the assessment of ploidy for all Columbia River White Sturgeon families reared at the Kootenay Sturgeon Hatchery. The standard protocol set by the Technical Working Group is that individuals found to be 12N will not be released into the wild (i.e., must be culled).

During the marking and tagging process, fish were sampled live using a diabetic syringe and collecting blood via the caudal vein. Collected blood samples were then dispensed into labelled heparinized vacutainer tubes, the tubes were organized into Styrofoam trays, bagged and then shipped on ice pack overnight via Air Canada cargo to the FFSBC fish health lab in Duncan, BC.

Once at the lab, capillary tubes are used to extract blood from the vacutainer. One drop of blood is added onto the inside of a sample vial lid and the lid then placed on the vial and gently inverting 2 to 3 times to mix. Each vial is prepared by combining 8 mL Isoton II diluent solution with 2 drops of Zapglobulin (Zapglobulin is a proprietary solution that lyses cell walls but leaves the nucleus walls intact). Each sample is then individually measured using a Z2 Coulter counter. The Coulter counter (Beckman Coulter Z2) has been used at FFSBC fish health lab since 2018 and is a much more efficient and accurate method to assessing ploidy compared to the historical blood smear method (see Fiske et al. 2019 for comparison of methods). Coulter counters operate on the same basic premise of blood smears using the size of cell nuclei to

determine ploidy status, but the counter allows 5,000-20,000+ nuclei from each sample to be measured. The FFSBC fish health lab has determined that the optimal detection range for ploidy samples from salmonids and sturgeon is an upper size (Tu) of 9 microns and the lower size (Ti) of 3 microns. Beckman Coulter's Z2 Accucomp software is used to collect the data (specifically the mode i.e., the most numerous nuclei size in that range) for each sample and to export it to MS Excel. An advanced Excel sort procedure is used to align sample numbers with the calculated mode of each sample. These modes are then compared to modes from samples of known ploidy status (e.g. sturgeon 8n and 12n).

2.4 Release Targets

The conservation aquaculture program has been adaptively managed in response to post-release monitoring results since its inception. Analysis of survival from previous stocking events has shown that juvenile survival improves with size-at-release and is expected to be approximately 86% in the Canadian portion of the Columbia River below HLK dam at a release size of 200g. Since the incorporation of wild-origin progeny, there has been a concerted effort to attain a minimum release size of 200g for fish released in the Columbia River below HLK and 300g for fish released above HLK into Arrow Lakes (Mid Columbia River) in order to maximize survival of these more genetically diverse groups. Further, wild-origin progeny from the Revelstoke spawning site are reared to even larger sizes (~700 g) in an attempt to further promote survival as progeny from wild adults residing in Arrow Reservoir have only recently been incorporated into the aquaculture program. The increased size target for Arrow Lakes is due to apparently low survival as very few hatchery-origin fish have been recovered to date, despite intensive sampling effort. Numbers of fish released into Arrow Lakes were higher earlier in the program when broodstock progeny were available in large numbers but the targets are now reliant on wild-origin collections which leads to annual variation and lower average release numbers. Conversely the target number of fish to be released in Canada below HLK dam has decreased over time (with knowledge of high survival rates) and is currently set at a maximum of 200. For the full retrospective time series on stocking targets, see Table 15 in: Mid and Lower

Columbia River White Sturgeon Conservation Aquaculture Comprehensive Report: Study Period 2001 to 2019 (BC Hydro 2019).

While the specific release number and size-at-release targets are largely driven by objectives to recover the Columbia White Sturgeon population to specified adult spawner abundance levels; there are also broad recovery objectives in regard to maintenance of genetic diversity. This objective is considered in how the specific wild origin spawn groups are segregated and released to specific locations. Release locations are selected with the goal of releasing progeny back to an accessible site as close as possible to the spawning site of their original capture. The mix of fish released to a location is set with the objective of including the full diversity of spawning groups from that source location and then using surplus to augment numbers at locations where sufficient embryos were not captured to fulfill the spawning target numbers. To date, Waneta has produced far more fish for release than other sites and is the only location where enough progeny are captured to satisfy the target for the lower Columbia. Therefore, generally all fish from sites other than Waneta are released back to their source location, and Waneta fish are used to make up the balance of the target number. All surplus are released into Arrow Lakes, following annual discussions with agencies and the technical working group.

3. Results

3.1 Captured embryos and larvae

Wild embryos and larvae have been captured annually since 1993 and incorporated into the hatchery program starting in 2014, however, capture rates across years can be variable. Waneta has been the most consistent capture location, and this was the case for the 2022 brood year (all larvae were from Waneta), but there was also 1 egg collected at Kinnaird and 18 eggs collected at Revelstoke. The number of eggs and larvae from each spawn group brought to Kootenay Sturgeon hatchery in 2022 is summarized in Table 3 (refer to Figure 1 for site locations) and totalled 527 larvae and 19 eggs. The large number of collection events, 19, necessitated that the initial capture groups be combined into larger rearing groups that remain consistent throughout the entire rearing process. The combined collection groups were always

from the same spawn location, and therefore a final group code combined a numerical rearing group and the first letter of the source spawning location. This approach allows for easy interpretation of the relative sequence and source location of each group and consistent throughout the rest of the report results. Ultimately, there were four groups from Waneta and one group from Revelstoke, and the one egg captured at Kinnaird did not survive.

Table 3. Wild captured eggs and larvae transferred to the Kootenay Sturgeon hatchery in 2022.

Brood Year	Check-In Date	Larvae	Eggs	Collection Site	Collection Group	Rearing Group	Group Code
2022	8-Jul-2022	52		Waneta	1	1	1W
2022	8-Jul-2022	3		Waneta	2	1	1W
2022	8-Jul-2022	245		Waneta	3	1	1W
2022	8-Jul-2022	29		Waneta	4	2	2W
2022	8-Jul-2022	2		Waneta	5	2	2W
2022	12-Jul-2022	1		Waneta	6	2	2W
2022	12-Jul-2022	1		Waneta	7	2	2W
2022	12-Jul-2022	2		Waneta	3	1	1W
2022	18-Jul-2022	25		Waneta	8	3	3W
2022	18-Jul-2022	63		Waneta	9	3	3W
2022	18-Jul-2022	1		Waneta	10	3	3W
2022	26-Jul-2022	64		Waneta	11	3	3W
2022	26-Jul-2022	2		Waneta	12	3	3W
2022	Did not survive		1	Kinnaird	13		K
2022	26-Jul-2022	18		Waneta	14	4	4W
2022	26-Jul-2022	16		Waneta	15	4	4W
2022	26-Jul-2022	2		Waneta	16	4	4W
2022	5-Aug-2022	1		Waneta	17	4	4W
2022	18-Aug-2022		12	Revelstoke	18	5	5R
2022	26-Aug-2022		6	Revelstoke	19	5	5R

3.2 Juvenile growth, survival

In-hatchery growth trends were documented at several points through the rearing year after the fish are received during summer. For the 2022 brood year fish, there were two bulk weight

sample events (a total weight taken and the number of fish counted to get average weight) on September 9th and December 2nd. Individual level data was recorded started with tagging and tissue sampling on February 22, and individuals were sampled in April, May and June for inventory, pre-release sampling and post release sampling of hold-overs.

The complete growth trajectory of each group from fall through to spring of the next year prior to release is documented in Figure 4. The size distribution of fish as measured during the pre-release sort showed the modal weight of fish to be near the 200g release target for lower Columbia. However, the distribution was right-hand skewed with significant variation in size, so that most fish were well in excess of the release target. Incredibly, one fish reached a size of 818g in less than a years time of being received in the hatchery (from group 3W3).

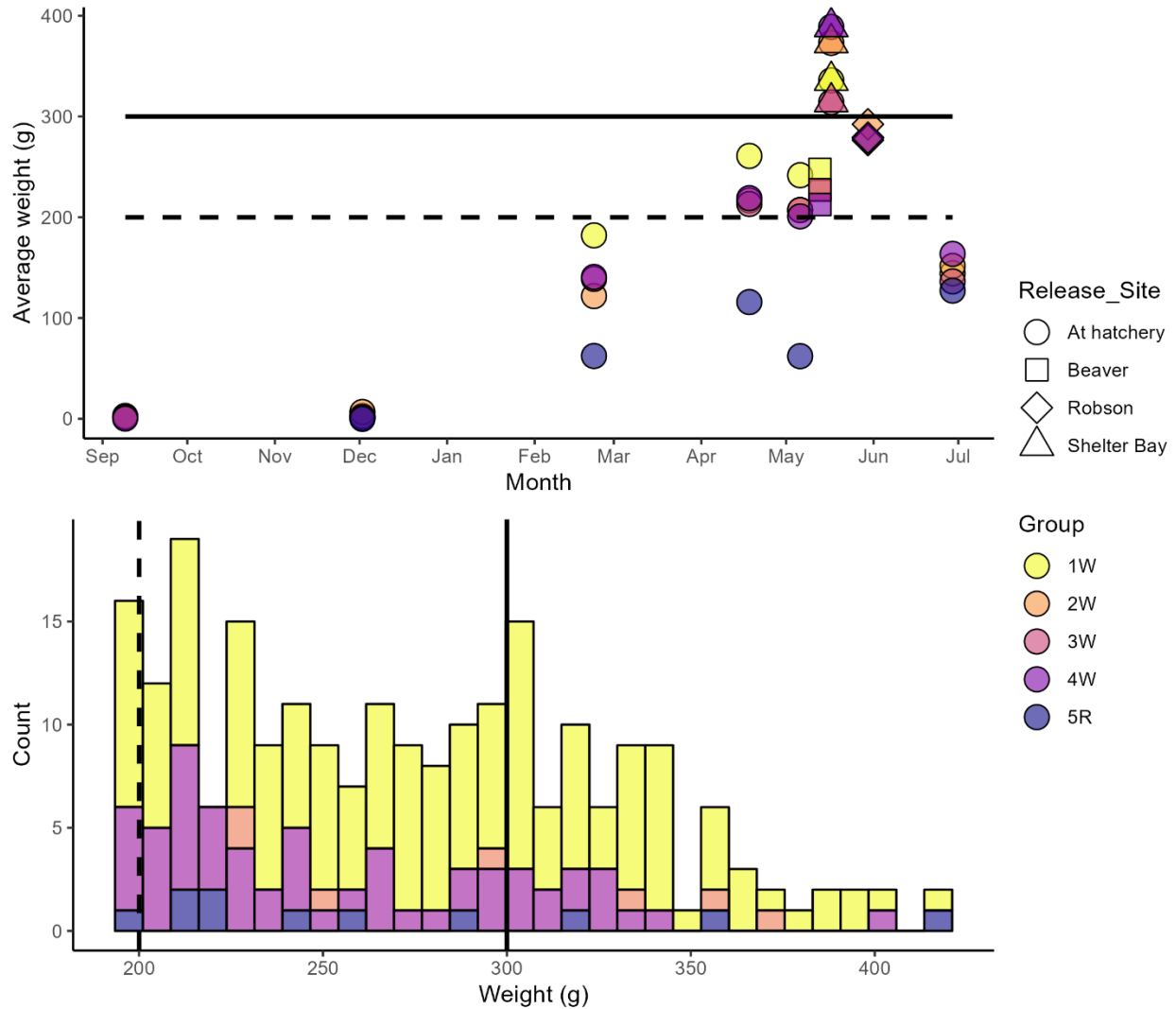


Figure 4. Growth trajectory over time (upper) and size distribution pre-release (lower) of all wild juvenile white sturgeon groups from 2022 brood year reared in Kootenay hatchery. Reference lines refer to minimum size-at-release targets for Columbia River (dashed) and Arrow Lakes reservoir (solid). Colour scheme in both panels match legend)

In the more recent years of wild origin progeny collection there has not been a culling of fish, and so it is possible to observe the overall survival and growth rates from received larvae to yearlings (or older) released. Table 4 summarizes the survival rates for each of the larvae and egg groups captured in 2022. For comparison the all-year average survival rate for larvae is 48%

and for eggs it is 21%. Overall, the combined survival of all groups from 2022 brood year to spring 2023 was 56%, which is above average.

Table 4. Numbers of wild larvae and eggs received at Kootenay Sturgeon Hatchery in 2022 and the resulting number of juveniles (yearlings or older) released or held for future release (at a larger size).

Brood Year	Group	Larvae	Eggs	Released	Held over	Survival rate
2022	1W	302		154	30	61%
2022	2W	33		7	8	45%
2022	3W	155		57	30	56%
2022	4W	37		11	10	59%
2022	5R		18	0	7	39%
2022	K		1	0	0	0%
2022	UNK	NA		NA	1	NA
2022	Total	546	19	229	86	56%

86 fish were held over for future release into Arrow lakes at a larger size, this included all of the seven-remaining fish of Revelstoke origin.

3.3 Juvenile sampling, marking and testing

The 2022 brood year fish were initially pit tagged and tissue sampled on February 22, 2023, thus individual level sampling information is available for that date as well as pre-release sorts (in May) and/or final inventory on June 29, 2023. The average sizes per group are captured in the Figure 4 above.

Autopolyploidy

Results from screening of hatchery reared fish (since brood year 2013) have demonstrated the 12N occurrence to be very low (Table 5), but fish originating from broodstock have significantly higher rates of occurrence than fish originating from wild captured embryos (BC Hydro 2019 CLBWORKS 24,25,34). In the 2022 captured larvae, 1 of the 317 fish (0.3%) sampled by the FFBC fish health lab in Duncan, BC was 12N. This fish was removed from the groups to be released.

Table 5. The proportion of hatchery reared fish testing positive for 12N ploidy.

Origin	Brood Year	Method	Sample	12N	Proportion 12N
Hatchery	2013	ImageJ	180	9	0.050
Hatchery	2014	ImageJ	1,693	124	0.073
Wild	2014	ImageJ	1,098	2	0.002
Wild	2015*	ImageJ	63	1	0.016
Wild	2016	ImageJ	1,209	1	0.001
Wild	2017	ImageJ	737	3	0.004
Wild	2018	Coulter	1,063	4	0.004
Wild	2018*	ImageJ	112	5	0.045
Wild	2018*	Coulter	112	7	0.063
Wild	2019	Coulter	302	0	0.000
Wild	2020	Coulter	710	10	0.014
Wild	2021	Coulter	519	6	0.012
Wild	2022	Coulter	317	1	0.003

* 2018 Paired re-sampling of same 112 individuals to compared methods.

* In 2015, an additional 2,154 wild origin sturgeon larvae captured in the USA were transferred to Kootenay Sturgeon hatchery and all 2,154 were confirmed as 8N using Image J method.

3.4 Juvenile releases

The release targets in numbers (200) and size targets (200g) were met for the Columbia River below HLK in 2023. The timing of release is generally focused on the month of May but hold back to a later time does occur to ensure the size targets, health and polyploidy testing are all achieved. In the spring of 2023 sturgeon were released on May 13th, 17th and 30th. The sturgeon released below HLK at Robson boat launch and Beaver Creek were primarily of 2022 brood year, but 11 (much larger) individuals from the 2021 brood year were also released at Beaver Creek. The size targets for release above HLK into the Arrow Lakes reaches are much larger (700g and over for any Revelstoke origin fish, and 300g or over for surplus lower Columbia origin fish) which requires additional rearing time. This was the case in 2023 releases with 2021 sturgeon making up most of the release (104) and averaging very large sizes (>600g). An additional 40 individuals from the 2022 brood year were released above HLK and averaged just over the 300g target at 336.15. Detailed information on the release location, collection site, collection group, the number released and average weight at release are all summarized in Table 6. All of this information is recoverable on recapture through pit tag identification and will

allow for analyses of potential effects of different collection site and/or release strategies on growth and survival in the future. Occasionally, the association to an original collection group (collection site and spawn event) for a fish is lost due to pit tag extrusion or combining groups (post tagging) for rearing. However, this usually does not prevent association of the new pit tag to the correct collection site, if all groups (spawn events) within the container originated from the same collection site. A total of 26 of the 2021 brood year fish had lost their original collection group information at time of release, but all were known to be of Waneta origin. This was an unusually high rate of tag loss, which was suspected to be due to less experienced staff participation (an adjustment was made after this observation and tag loss was low again in 2022). Almost half (71) of the fish released above HLK were originally collected from Revelstoke in 2021, and had the largest average mass (707g) of any group.

In 2022 a total of 121 large (>300g) sturgeon were released above HLK into the Arrow Lakes section of the river at Revelstoke or Shelter Bay, and all of those fish were originally collected from Waneta. Exactly 200 individuals (>200g) were released below HLK, evenly split between Beaver Creek and Robson Boat launch release locations. All of the fish released below HLK were originally captured at Waneta and distributed among 6 different spawn groups.

Table 6. All White Sturgeon releases to the Columbia River in 2023.

Brood Year	Release date	Release Site	Collection Site	Group	Number Released	Average Wt (g)
2021	May 17, 2023	Centennial Park	Waneta	3W3	1	370.0
2021	May 17, 2023	Centennial Park	Waneta	4W4	2	675.0
2021	May 17, 2023	Centennial Park	Waneta	5W5	1	610.0
2021	May 17, 2023	Centennial Park	Waneta	Unknown	18	601.1
2021	May 17, 2023	Centennial Park	Revelstoke	10R1	1	580.0
2021	May 17, 2023	Centennial Park	Total		30	597.3
2021	May 17, 2023	Shelter Bay	Waneta	1W1	1	360.0
2021	May 17, 2023	Shelter Bay	Waneta	Unknown	3	620.0
2021	May 17, 2023	Shelter Bay	Revelstoke	10R1	70	716.1
2021	May 17, 2023	Shelter Bay	BY 2021 Total		74	707.4
2022	May 17, 2023	Shelter Bay	Waneta	1W	31	335.8
2022	May 17, 2023	Shelter Bay	Waneta	2W	1	373.0
2022	May 17, 2023	Shelter Bay	Waneta	3W	6	314.3
2022	May 17, 2023	Shelter Bay	Waneta	4W	2	389.0
2022	May 17, 2023	Shelter Bay	BY 2022 Total		40	336.2
Total above HLK					144	
2021	May 13, 2023	Beaver Creek	Waneta	1W1	2	530.0
2021	May 13, 2023	Beaver Creek	Waneta	4W4	3	686.7
2021	May 13, 2023	Beaver Creek	Waneta	5W5	1	790.0
2021	May 13, 2023	Beaver Creek	Waneta	Unknown	5	439.0
2021	May 13, 2023	Beaver	BY 2021 Total		11	555.0
2022	May 13, 2023	Beaver Creek	Waneta	1W	54	247.7
2022	May 13, 2023	Beaver Creek	Waneta	2W	1	227.0
2022	May 13, 2023	Beaver Creek	Waneta	3W	29	227.3
2022	May 13, 2023	Beaver Creek	Waneta	4W	5	212.8
2022	May 13, 2023	Beaver	BY 2022 Total		89	238.9
2022	May 30, 2023	Robson	Waneta	1W	69	279.4
2022	May 30, 2023	Robson	Waneta	2W	5	292.2
2022	May 30, 2023	Robson	Waneta	3W	22	276.5
2022	May 30, 2023	Robson	Waneta	4W	4	278.5
2022	May 30, 2023	Robson	Total		100	279.4
Total below HLK					200	
Grand total released in 2023					344	396.7

- Refer to Table 3 in BC Hydro 2022 report for 2021 BY group code description, and this report (Table 3) for 2022 brood year group code description.

Over 180,000 sturgeon have been released as yearlings or older since the inception of the conservation aquaculture program (Table 7). One key finding gleaned from the historical releases is that survival is size dependent (BC Hydro CLBMON-29 2016), and a trend towards larger release size is notable in the stocking records (Figure 5). The fish released in 2022 achieved an average size well in excess of the target size-at-release (Figure 5). Although as seen in Table 6 a portion of the release above HLK included 2022 brood year sturgeon that averaged slightly below 300g at 272g. At present, the large release size targets are seen as an important measure to ensure survival of the more genetically diverse wild origin progeny.

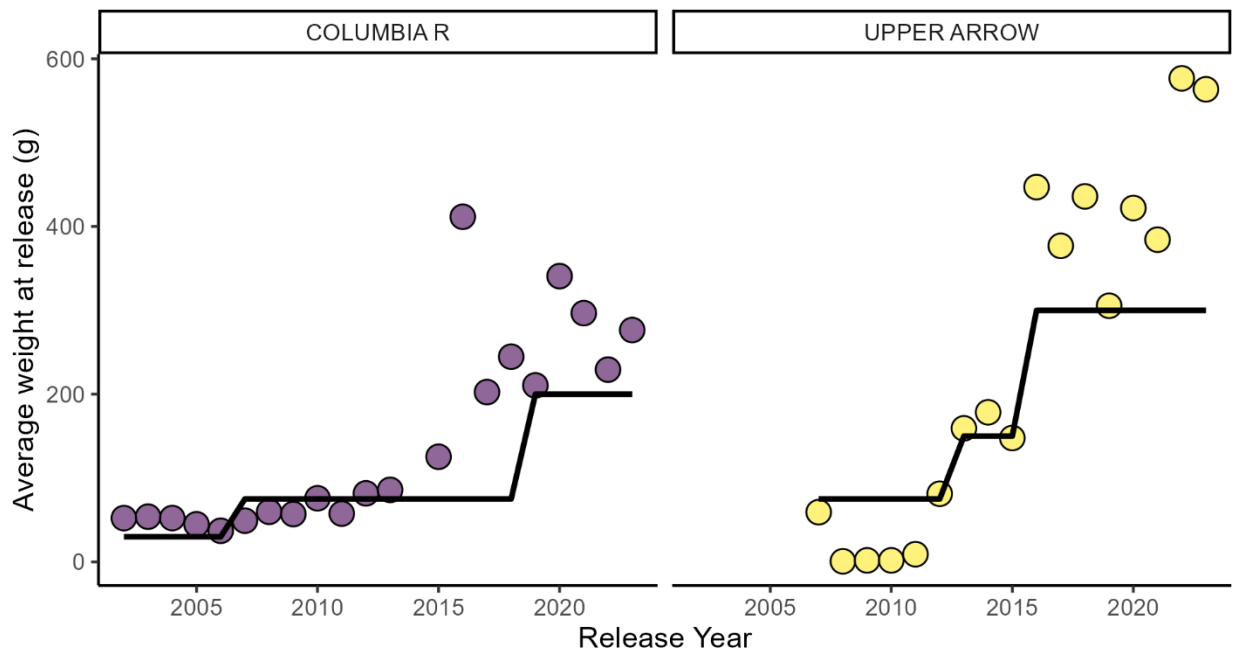


Figure 5. Trends in release size (g) over the full span of the aquaculture program to spring 2023. Each point is the average size for the release year across groups. Panels are labelled by release location: Columbia R = All release sites below HLK, and Upper Arrow = All releases size above HLK. Points appearing as 0g, were larvae experimentally released into Upper Arrow. The solid back lines depict the minimum size-at-release targets, which changed for both areas over time.

Table 7. All releases of brood and wild origin progeny reared at Kootenay Sturgeon Hatchery and released into the Columbia River. Columbia River zones refer to five 11.2 km sections between Hugh Keenleyside Dam and the USA border.

Source	Release Year	Columbia Zone of Release					Arrow	USA	Total Releases	
		1	2	3	4	5				
Brood	2002	4,407				4,264			8,671	
Spawn	2003	5,724				6,079			11,803	
	2004	4,828				4,867		1,882	11,577	
	2005	8,996				8,881		3,755	21,632	
	2006	7,267				7,603			14,870	
	2007	6,237				5,915	4,206		16,358	
	2008	3,216				3,233	6,534		12,983	
	2009	3,738				400	8,118		12,256	
	2010	3,458	15	14	15	515	9,575		13,592	
	2011	2,500			1,000	510	8,078	2,621	14,709	
	2012	3,189			1,000		6,567		10,756	
	2013	2,293			875		869	5,944	9,981	
	2014	1,200			300		301	6,017	7,818	
	2015	1,600			600		600	3,288	6,088	
	Wild Capture	2015					1,095			1,095
		2016					76	1,324		1,400
2017						800	1,589		2,389	
2018		378				229	975		1,582	
2019		100				100	541		741	
2020		100				100	212		412	
2021		100				99	285		484	
2022		100				100	121		321	
2023		100				100	144		344	
Total		59,531	15	3,789	15	46,736	63,518	8,258	181,862	

*Not yet released

4. Discussion

The Upper Columbia White Sturgeon Recovery Initiative has accomplished a vast amount of work towards numerous aspects of recovery of this population (e.g. see Upper Columbia White Sturgeon Recovery Initiative Operational Plan updated annually by the technical working group). The conservation aquaculture program has been a vital component of the initiative and

the successful establishment of hatchery-origin juveniles in the Columbia River has prevented extirpation of the population (Crossman et al. 2023). The present abundance of sturgeon in the upper Columbia is estimated to be 5 to 10 times greater than without intervention for Canada and USA respectively. Moreover, one of the key recovery plan objectives of 2,500 reproductive adults in Canada appears to be numerically achievable with the current stocking program given current estimates of survival and maturation rates of hatchery fish. Albeit it is still not certain how higher densities of fish and decreased growth rates may slow maturation rates and delay achieving the “reproductive” status.

The 2022-2023 sturgeon aquaculture program has once again been very successful. All targets for release numbers and size-at-release were met, and survival rates of wild-origin progeny during rearing at the hatchery were above the long-term average. Survival of larvae at the time of transfer to the hatchery was slightly above average at 58% and the eggs collected from Revelstoke had higher survival in the hatchery (37%) compared to prior years. While eggs collected at Revelstoke survived better, the spawning site remains challenging due to significant sand and debris encountered during sampling which can negatively affect survival. In 2023, the streamside trailer may be relocated to Revelstoke once the lower section wild progeny collection is complete. The rationale would be to increase survival of embryos that are collected by reducing the amount of handling required during this critical early development period.

The change to wild-origin progeny is assumed to have significantly increased the genetic diversity of the released juveniles, though this needs to be further confirmed through genetic analyses. A comprehensive program is underway to describe genetic diversity of wild-origin progeny released from this program since 2014 and results are expected in 2024. A key objective of the genetic work will be to examine genetic diversity between spawning events within year classes as well as between stocking years. It is expected that results from this genetic work will greatly inform next steps of the aquaculture program and release strategies. Most of the wild progeny continue to be captured only at Waneta, but effort to represent as many spawning locations as possible is ongoing. For example, a spawning habitat restoration

project was completed at the ALH site in 2023 with the objective of improving conditions for early life stage development and survival. The release strategy for individual spawn groups within the Waneta site has varied slightly over the years. All of the 2021 brood year fish were of Waneta origin, so there was no need to allocate fish to specific release groups in order to match capture location with proximity to release location. The specific Waneta fish selected for release above HLK were selected based on size to meet the 300g target. Allocation to the two lower locations attempted an approximately equal distribution for each group across locations to ensure each spawning event was represented in each release location. In future years, it may be worth considering adaptive management of release group-location strategies based on any updated understanding of the best approach for long term genetic diversity.

Autopolyploidy was a minor issue in the 2022 brood year class with only 0.3% confirmed 12N. This is towards the lower range of historical (wild origin larvae) values, despite using the more accurate Coulter counter method which has a higher detection probability than the historical blood smear methods.

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