

STANISLAUS RIVER
ROTARY SCREW TRAP PROTOCOL

June 2016

**STANISLAUS RIVER ROTARY SCREW TRAP PROTOCOL
TABLE OF CONTENTS**

INTRODUCTION	4
SAFETY	4
COMPLETING DATA SHEETS	6
TRAP STATUS, ENVIRONMENTAL DATA, AND FISH CATCH DATA COLLECTION	8
Trap Status and Environmental Data	8
<i>Cleaning the Trap</i>	9
Fish Catch Data Collection	10
<i>Setting Up and Maintaining Buckets, Insulated Coolers, and Live Carts</i>	10
<i>Collecting Fish</i>	10
<i>Sub-Sampling Protocol</i>	12
<i>Processing Fish</i>	12
<i>Anesthetizing Salmon That Are Measured</i>	14
<i>Collection of Fin Clips</i>	15
Field Quality Check	15
Trap Maintenance	16
MARK/RECAPTURE TRIALS	16
COMPUTER DATA ENTRY AND MANAGEMENT	19
Data Entry	19
QA/QC Procedure.....	19
APPENDIX A: Field Data Sheets.....	21
Trap Visit Data Sheet.....	21
Unmarked Chinook Salmon Data Sheet.	22

Unmarked Steelhead Data Sheet.....	23
Spring-run and Winter-run LAD Chinook Salmon Fin Clip Data Sheet.....	24
Photonically Marked Chinook Salmon Data Sheet.	25
Recaptured Chinook Salmon Data Sheet.....	26
Adipose Fin Clipped Salmonid Data Sheet.	27
Other Species Data Sheet.....	28
APPENDIX B: Chinook Salmon and Steelhead Life Stages	29
APPENDIX C: Trap Visit Data Sheet Terminology	31
APPENDIX D: Equipment Lists	36
Catch Visit Equipment List.....	36
Bismark Brown Equipment List	36
Anesthetizing Equipment List.....	36
Elastomer Dye Equipment List.....	36
APPENDIX E: Fish Species List for The Stanislaus River.....	38
APPENDIX F: Key to Juvenile Chinook Salmon and Steelhead	40
APPENDIX G: Process for Staining Chinook Salmon With Bismark Brown Dye	41
APPENDIX H: Process for Marking Chinook Salmon With Elastomer Dye	42
APPENDIX I: Operational Procedure for Night Operations	45

INTRODUCTION

Monitoring data can provide the foundation for successful management programs if data are collected in a systematic, consistent, and comprehensive manner. The monitoring of the abundance/production of juvenile salmonids on the Stanislaus River is important because large sums of funding are spent restoring aquatic habitat in an effort to increase the number of juvenile Chinook salmon (*Oncorhynchus tshawytscha*) and steelhead (*Oncorhynchus mykiss*) in that watershed. Rotary screw traps (RSTs) are one of the most important tools for monitoring juvenile salmonids. When data acquired with these tools is collected in conjunction with other monitoring data, there is a substantially improved ability to track the status of those salmonids and assess their response to past management activities. That data in turn can be used to adaptively manage future restoration projects so that they are more successful.

The objective of this document is to ensure that RST data from the Stanislaus River in California's San Joaquin Valley is collected in a safe, systematic, consistent, and comprehensive manner. To address this objective, this protocol provides detailed descriptions for operating and maintaining RSTs, collecting and processing fish, collecting environmental data, conducting trap efficiency tests, and entering data into a RST "platform" developed by the U.S. Fish and Wildlife Service's Comprehensive Assessment and Monitoring Program (CAMP).

The Stanislaus River RSTs are located at Caswell State Park, approximately 5 miles southeast of the town of Ripon in San Joaquin County, California. A total of two 8 foot RSTs will be operated each year. Fall-run Chinook salmon and steelhead/rainbow trout are the most likely salmonids to be captured in the Stanislaus River RSTs.

The CAMP has developed a general protocol for conducting RST activities. That document can be found at https://www.fws.gov/cno/fisheries/CAMP/Documents-Reports/Documents/2008_draft_CAMP_Rotary_Screw_Trap_Protocol.pdf. The general guidance in that document should serve as a companion to the more detailed guidance in this document.

Employees who are engaged in servicing the RSTs on the Stanislaus River are responsible for having a firm understanding of all the information covered within this protocol.

SAFETY

Safety should always be the field biologist's primary concern. Never perform a task if it cannot be performed safely. A minimum of two crew members are required to service RSTs at any given time. At least one biologist must have a working cell phone when in the field. RST activities occur in an environment where field biologists can be seriously injured by rotating mechanical parts, crushed between a boat and a trap, or drown if they do not wear personal protective gear. The Stanislaus River RSTs operations have the potential to seriously injure members of the public who use the river for recreational purposes if those individuals become caught in the trap. The proximity of the Stanislaus River RST site to the cities of Modesto, Ripon, and Stockton also creates the potential that field biologists servicing the traps could be

exposed to discarded hypodermic needles or pathogens that may be present in the river. These situations, in combination, therefore demand that field biologists servicing the Stanislaus River RSTs pay attention to, and implement, measures that are designed to ensure their safety and the safety of the public. Because there may be a considerable amount of interaction between the public and the RST field crew, crew members should display a high level of professionalism when they are in the field.

Crew members are required to wear their personal flotation devices at all times when they are on a boat or a trap.

Proper communication is essential when operating a motor vessel. When piloting a boat, the boat captain should make sure their intentions are clear to boat passengers. The captain should give clear warnings before they are about to engage or disengage the throttle, or make a turn. This will give boat passengers proper time to sit down and become stable. The small boat used on this project is very unstable when standing or kneeling, and may be easy to fall off when an unexpected move is made.

Great care should always be taken when working on a RST. Be cautious to always keep hands, loose clothing, and other items away from the cone, shaft and other moving parts during trap operations. Be cautious when moving around on a trap because numerous hazards exist, e.g., the winch, cleats, cables, frayed cable, etc. Familiarize yourself with these hazards and use caution when moving about. Never move across the number one crossbeam (in front of the trapping cone) when the trap is fishing. If traps have been modified by installing catwalks in the front of the trap, biologists should use caution when they are on the catwalk and avoid the potential that they could intentionally or accidentally end up inside the trap cone.

Always be aware of other crewmember locations and activities on the trap. Keep alert for boat traffic and boat wake, and during high flow conditions watch for large debris that may collide with the RST or boat.

When pulling the boat up to a trap, all crew members should give this part of the operation their full attention. Do not place any part of your body in between the boat and a trap during an approach or while moored to a trap; together, the boat and trap can crush and cause severe injury to your body. If available, put the boat fenders over the side before pulling alongside a trap. The boat operator should drive the boat slowly when approaching a trap. The boat operator should carefully maneuver the boat alongside the trap pontoon so a crewmember can step, not jump, from the boat to the pontoon. Once someone is safely on the trap and the boat has been secured to the trap, the boat operator can shut down the boat motor. The crew should make sure that the fenders are adjusted properly to prevent the boat and trap from rubbing together and that the boat is securely tied to the trap. The boat should not be operated directly in front of the trap; if the boat loses power, the current will push the boat into a trap cone and possibly damage and/or sink the boat. Assume the jet boat can lose the ability to propel itself when debris gets caught into the intake of the jet pump, and plan accordingly.

Transferring gear or persons between the boat and trap should occur with the full knowledge of all crew members. One person should be on the boat and another on the dock/trap and gear

should be transferred from crew member to crew member. Be very careful when stepping on or off the trap or walking on the trap. The pontoons and live-box lid may be slippery any time they are wet or biologist's boots/shoes are wet, but they can be especially slippery due to frost or ice in the winter or algal growth in the spring and summer.

Check the winch cable and mooring cables for fraying. Use caution when handling cables to avoid punctures to hands from frayed cable. Check the following for damage: carabineers, quick links, cleats, and eyes. If damage is found, notify the entire field crew of the damage and take steps to promptly repair that damage. When raising or lowering the trap cone or live-box door, all crew members should know each other's whereabouts on or off the trap and everyone should be aware this procedure is taking place. One person should observe the trap cone while another person is raising or lowering the cone. Remember that it can be difficult to see the front of the cone when operating the winch. Crew members should look to make sure trap components are functioning properly, there are no obstructions in the pulleys and the #2 beam, collars are secure, debris has not accumulated on the trap components, winch cables are not damaged, etc. The person initiating a procedure should make it known by saying aloud that the trapping cone or live-box door is about to be moved, and all others should acknowledge they are aware of what is about to take place. When the trapping cone is being lowered, keep toes clear of the cone to prevent them from being crushed between the #2 crossbeam and the pontoon. Always secure the live-box door if it is in an open position to preclude the potential of being hit by the live-box door.

The crew should observe weather and river conditions prior to their day in the field. Utilize the guidance plots on the Department of Water Resources CDEC website to see if a forecasted increase of river rise and flow may occur when out in the field. On days when the river may rise, the crew must go to the field with waders and vest PFD's. Such days tend to coincide with heavier and larger debris that may get caught on the traps.

COMPLETING DATA SHEETS

Properly completing data sheets is critical to project success; this aspect applies to every data sheet used during the Stanislaus River RST project. Data sheets should be clear, legible, and contain all information needed to accurately interpret data at a later date, i.e., a data recorder should ask themselves if someone could accurately and completely infer what data were collected five years after the form was filled out. If there is more than one data sheet for a particular site, make sure they are labeled appropriately (e.g., site name, page 1 of 2, etc.). Make all information clear enough so someone not familiar with the field activities can interpret the data accurately (i.e., use standard abbreviations, no omitted data). There should never be any empty spaces for relevant data on a data sheet. If data are not taken, draw a line through the appropriate box and write a short explanation in the "Notes" section of the Trap Visit data sheet.

Additional comments regarding any variations in procedure, notable field conditions or other pertinent information should be recorded on the Trap Visit data sheet, e.g., any river conditions affecting trap operation or changes in a trap's deployed position should be noted.

Use the following conventions when filling out data sheets:

1. A data sheet will be filled out every time a trap is set and every time a trap is checked. If a site was sampled and no fish of any kind were caught, fill in all of the relevant fields on the data sheet with trap check and environmental data, and put a note on the data sheet explicitly stating no fish of any species were caught. If no salmonids were caught, but other fish species were captured, that should be explicitly noted on the data sheet. The occurrence of situations with no catch data is critical because it is used in the calculations of juvenile salmon production estimates.
2. If there were logistical or operational conditions that occurred and resulted in an atypical sampling period, completely and clearly explain what the conditions were in the “Notes” section on the data sheet. Keep explanations professional and organized for clarity.
3. Use a pencil, and your best and clearest non-cursive handwriting.
4. Organize the data sheet so data for the same fish species is recorded together. Look at the catch before data is recorded and leave ample space to group data for each species. Use additional sheets as necessary to assure clarity of information.
5. Completely fill out each portion of a data sheet. If a section of the data sheet is left blank, draw a line through it and explain why it wasn’t filled out in the “Notes” section of the data sheet.
6. Corrections can be made in the field by erasing a mistake if the sheet is dry, or putting a line through the mistake and clearly writing correct information nearby.
7. Never estimate data, i.e., record measured values only. If a value cannot be measured, put a line in the box where that value would be recorded and provide an explanation in the “Notes” section of the data sheet.
8. The lengths of dead fish should be recorded separately from live fish on the data sheet, regardless of whether the fish is a salmonid or not, and their mortality status must be noted.
9. Each fish that is measured for fork length and weight should be tallied separately on the data sheet from other fish that were only enumerated.
10. The presence and number of marked juvenile Chinook salmon that are recaptured should be recorded on the Trap Visit data sheet in the special section reserved for those fish. If more than one kind of mark combination is observed in the recaptured fish on a given day, take care to segregate the “plus count” of the number of fish with each kind of mark.
11. Plus counted fish need to be clearly labeled as “Live unmeasured” or “Morts unmeasured.” Those numbers should then be totaled on the Trap Visit data sheet.

TRAP STATUS, ENVIRONMENTAL DATA, AND FISH CATCH DATA COLLECTION

The procedures described below are presented in the sequential order that would be implemented in the field. This work flow has been designed to make field activities as simple, efficient, and complete as possible.

When traps are deployed in a “cone-down” position and are actively fishing, they will be checked at least once every 24 hours, and more frequently on days when there are high river discharge levels, heavy debris loads, or the capture of thousands of salmon could adversely affect the health of fish sequestered in RST live-boxes. On days when traps will not be checked within a 24-hour period, trap cones will be raised out of the water and placed in a “cone-up” position to preclude fish capture. On days when only one trap check occurs, there will be an effort to arrive at the trap between 8 AM and 10 AM.

Trap Status and Environmental Data

When the field crew initially arrives at the trap, they should inspect the trap for any signs of trespass or damage before they exit the boat and step on the RST. They should also examine the cone depth gage on the trap and record the cone depth on the Trap Visit data sheet. If no signs of trespass or damage are found, the crew can board the trap and collect and record data on the Trap Visit data sheet that characterizes the operational status of the trap and environmental parameters at the time of the trap visit. A separate data sheet should be prepared for each trap cone. The following data should be recorded on each Trap Visit data sheet:

- Trap visit date and time.
- Subsite code.
- Recorder/crew.
- Weather (including air temperature).
- Visit type ID.
- Trap functioning ID.
- Cone depth (needs to be documented before staff board the trap).
- Total revolutions since last trap check.
- Water temperature.
- Water dissolved oxygen.
- Water turbidity.
- Water velocity.
- Before cone cleaning revolutions (RPM).
- After cone cleaning revolutions (RPM).
- Dominant debris type and volume.

Refer to Appendix C for instructions on how to collect and record data pertaining to the trap status and environmental parameters.

Water temperature data loggers manufactured by the Onset Computer Company will be positioned inside the RST live-boxes to monitor water temperatures on a continuous basis.

Those data loggers should be downloaded and archived at weekly intervals, and the data from those loggers should be reviewed to detect the presence of water temperatures that may adversely affect salmonids.

Cleaning the Trap

Before all the fish have been removed from a RST during a day time check, the trap should be cleaned so it can operate as efficiently as possible. In general, trap cones are not cleaned and the counter is not reset during evening checks.

Never, under any circumstance, crawl or climb inside the RST cone while the trap is on the river.

Never remove debris from a trap cone or shaft that is rotating, i.e., always lock the cone in a non-rotating configuration before initiating the cleaning process. Remove any large debris from the front of the trap, clean the rear live-box screen with a scrub brush and rinse the debris off as best as possible. Under no circumstances should the live-box screen be removed unless sampling has been terminated. If the live-box screen needs to be removed, all fish and debris should be removed first and a new time should be recorded for Visit Time2 field on the data sheet. Sweep silt/debris out of the live-box, rinse debris off the trap deck, and check/clean pontoons for algae growth. If a gas-powered trash pump is used to wash algae off the trap cone, place it where it cannot come into contact with anything sensitive to heat; the exhaust and engine get very hot and can melt or burn things.

If a trap cone needs to be raised or lowered for cleaning purposes (or if the trap needs to be moved on or off the river), always be cautious when using the winch crank. When raising the cone, keep a hand on the winch crank handle and make sure the latch is caught in the gear securely. Latches tend to wear and if not secure the winch handle may spin quickly and cause injury. If the trapping cone needs to be lowered, one person will lower the trapping cone, while another guides the A frame. Be sure both individuals are aware of the other person's actions and coordinate their respective activities. The person on the winch should announce that the trapping cone is coming down while the other person should watch for problems as the trapping cone lowers. Once the trapping cone begins to spin, note what time the trap begins fishing and record the number of RPMs the trap makes in 60 seconds. Do this three times and calculate an average RPM for the 3 recordings in the "Before Cleaning RPM" field on the Trap Visit data sheet. The following days after the traps have been fishing in the water, will begin with cleaning the cones and taking 3 recordings of RPMs the trap makes in 60 seconds.

After the trap has been cleaned and is rotating in the water, the rotation rate of the cone should be assessed again and the RPMs recorded in the, "After Cleaning RPM" field on the Trap Visit data sheet.

Fish Catch Data Collection

Weather and river conditions will dictate the order of the trap cleaning and fish processing activities. When the weather is cool and the potential that temperatures could be a stress factor for fish is minimal, trap maintenance activities should occur first, and then fish get processed. This approach maximizes the amount of sampling to the fullest extent possible. If the weather is warm or fish could be adversely affected if they are held for a substantial period of time, the fish should be processed first, and then trap maintenance activities should occur. The text below assumes that trap maintenance activities will occur first since weather and river conditions will most commonly be benign.

If at all possible, the trap cone should not be raised before the fish are removed from trap's live-box. Raising the trapping cone creates a gap through which fish can escape, so it is best to clear the live-box while the trap cone is fully submerged in the water.

Setting Up and Maintaining Buckets, Insulated Coolers, and Live Carts

Prepare to remove fish from the live-box by filling buckets with river water. Live carts or insulated coolers will also be prepared for fish sequestration if more than 500 fish are observed in the live-boxes when they are first checked. When fish are being held in buckets, place the buckets in the shade and place lids on top of the buckets when they are unattended.

Use the dissolved oxygen (DO) meter to periodically check the water temperature and dissolved oxygen levels in the containers used to sequester fish. Make sure the difference in water temperatures and dissolved oxygen levels between the buckets/coolers and river does not exceed a 2° Celsius or 7–10 milligrams/liter difference, respectively. Add fresh river water to the buckets and coolers if they become too warm or experience depleted dissolved oxygen levels. If necessary, use an aerator to help maintain DO levels that are equal to, or greater than, the river water. Frozen water bottles may be added to containers used to hold fish on warmer days to help maintain cool water temperatures.

Collecting Fish

Begin the process of clearing the trap live-box by scooping larger debris out of the live-box with a pitchfork, making sure to search that debris for fish. If fish are found with the large debris, remove those fish and place them in a container with fresh water. As fish are collected, every fish is retained, regardless of whether it is dead or alive. Using 5 gallon buckets, quantify and record the amount of debris in the Debris Volume field on the Trap Visit data sheet, then jettison that debris behind the trap. Also note the type of debris caught and record under “Dominant Debris Type” on the Trap Visit datasheet. Collect man-made trash and dispose of it properly.

After a majority of debris has been removed from the live-box, use a large net to scoop smaller debris and fish out of the live-box. Scoop no more than 1/2 a net of fish/debris at a time, and gently empty the nets and place the contents in an 18-gallon tub of water. During sunny days the livebox deck can become quite hot; therefore, extra care should be taken to cool the deck with river water before emptying net contents onto the deck if such is necessary. Placing debris and

fish in an 18-gallon tub of water is the preferred, less stressful method of handling the fish. When removing fish from the live-box, be careful not to smash fish between the rim of the dip net and the wall of the live-box. The live-box corners are typically where fish get killed. If feasible, chase fish out of the live-box corners before attempting to scoop them with the net.

Carefully sort through the debris using a stick, salad tongs, or other probe, **DO NOT use your hands**; hypodermic needles or other sharp objects could be present.

It is important to sort the fish from debris as quickly as possible without overlooking any fish. Carefully find and remove all fish, remembering that some will be very small. As fish are removed from the debris, sort them and place them in separate containers as follows:

- a) All steelhead
- b) Marked fall-run Chinook salmon.
- c) Unmarked fall-run Chinook salmon.
- d) Non-salmonid species, including larger piscivorous fish that could potentially feed on smaller fish as fish are held and processed.

Field biologists who have received adequate training should be able to recognize juvenile steelhead based on morphological features; Appendix F provides reference material that can be used to disseminate between steelhead and Chinook salmon. **The presence of marked fall-run Chinook salmon can be assessed by looking for fish with a colored mark or Bismark brown dye; it is especially important to examine every salmon to see if they have these marks since their detection will drastically affect the ability to develop salmon production estimates.** The presence of non-fall-run Chinook salmon can be ascertained by referring to the length-at-date charts developed by Fisher and Greene, i.e., salmon that have a fork length that is outside the length range for a given sample day could therefore be spring-, winter-, or late-fall-run Chinook salmon.

Make sure fish are not over-crowded (i.e., < 25 smolts or < 50 fry per bucket; 100-150 individuals per standard-size cooler). If fish exhibit strange behavior, transfer them to another bucket/cooler to replenish oxygen and gently lower water temperatures. Fish that flare their gills or starting to gather along the surface of the water is a sign of reduced oxygen levels. With a large quantity of fish in the bucket, this can happen very quickly, especially when the weather is warm.

At the moment the live-box is clear of fish and smaller debris, two measurements must be taken and recorded immediately on the Trap Visit data sheet: (1) enter the time in the Visit Time 2 field and (2) record the Total Revolutions Since Last Trap Check on the lever actuated mechanical counter. These two values represent the end of the current sample and the start of the next sample. If the trap is stopped by debris, record the Total Number of Revolutions on the counter and explain the circumstances in the “Notes” section of the Trap Visit data sheet. If the trap cannot immediately be returned to a mode where it can collect fish and instead needs to be taken out of service for some period of time for cleaning/repair, a time will be entered in

the Visit Time2 field which reflects what time the trap was put back in service. On most days when the trap essentially runs on a continuous basis, biologists will only be entering a time in the Visit Time2 field on the data sheet. The Visit Time field should be filled out from the previous day's Visit Time 2 field. The Visit Time 2 field reflects the time that the trap was completely cleared of fish and debris for one day, and represents the start of the next sampling period that ends on the following day.

Sub-Sampling Protocol

The following subsampling procedure only pertains to the selection of Chinook salmon that are assessed for fork length, weights, and life stage. All other fish are referred to as "plus counts." These procedures do not relate to the determination of how many salmon of any run were caught on a given day because the procedure for quantifying the number of salmon caught always involves the counting of each individual fish. It is important to ensure that the subsampling has a true random sample to ensure no bias in the size and race distribution, and ensure that it is an accurate representation of what was captured as a whole.

Processing Fish

After the trap has been cleaned and resumes fishing, the collected fish should be processed in the following order: (1) all steelhead, (2) unmarked fall-run Chinook salmon, (3) adipose fin clipped/hatchery origin fall-run Chinook salmon, (4) efficiency marked/recaptured Chinook salmon, and (5) non-salmonid species. The table below identifies how each group of fish should be processed, and the data from those individuals should be recorded on their respective data sheets. Depending on the fish category, fish will be counted, evaluated for species, measured for fork length, measured for weight, assessed for life stage, checked for mortality, and/or evaluated for salmon run type and have a fin clip taken for salmon run assessment (for genetic verification if LAD is outside of the fall-run criteria).

Sampling Strategy For Different Fish Species That Are Collected During Rotary Screw Trap Operations On The Stanislaus River				
Steelhead	Fall-run Chinook Salmon	Ad-Clipped/Hatchery Fall-Run Chinook Salmon	Efficiency Test Marked Chinook Salmon	Non-Salmonid Species
Each Individual is:				
<ul style="list-style-type: none"> • Counted • Assessed for life stage • Measure up to 100 randomly selected for fork lengths • The first 25 of individuals that are ≥ 40 mm will be randomly selected for weights • If more than 100, “plus count” remainder 	<ul style="list-style-type: none"> • Counted • Assessed for life stage • Measure up to 100 for fork lengths • No weight taken • If more than 100, “plus count” remainder 	<ul style="list-style-type: none"> • Counted • Assessed for life stage • Measure up to 100 for fork lengths • No weight taken • If more than 100, “plus count” remainder 	<ul style="list-style-type: none"> • Counted • Assessed for life stage • Measure up to 100 randomly selected for fork lengths • The first 25 of individuals that are ≥ 40 mm will be randomly selected for weights • If more than 100, “plus count” remainder 	<ul style="list-style-type: none"> • Counted • Assessed for life stage (adult/juvenile) • Measure up to 50 of each species for fork length • No weight taken • If more than 50 of one species, “plus count” remainder

Fish species identification can be accomplished using various keys that include fish found in the Central Valley; Appendix E provides a species list for the fish that are likely to be caught in the Stanislaus River based on previous RST work, and the appendix identifies three references that can be used to identify fish species in California’s Central Valley. If a fish is collected and biologists are not sure of the species identification of that individual, several close up pictures of that fish should be taken with a digital camera. Biologists should then refer to the species list and references in Appendix E in an effort to determine the species ID. If a fish cannot be identified to species, then the data sheet should reflect a more general taxonomic name for that individual. This situation may be especially true for various groups that include lampreys, Cottids (sculpins), Centrarchids (bass and sunfish), Catastomids (suckers) and Cyprinids (minnows). Individuals pertaining to these groups should receive special scrutiny in an effort to make taxonomic assignments as accurate as possible.

ALWAYS check juvenile Chinook salmon for marks that could have been applied as part of the mark-recapture trap efficiency trials described below.

The collection for data pertaining to salmon life stage, fish weight and fork length is relatively straightforward. Salmon life stage will be assessed according to the morphological features described and illustrated in Appendix B. All fish length measurements for fish species with a forked tail will be measured for fork length to the nearest 1.0 millimeter. For species without a

forked tail (i.e., lamprey, sculpins, mosquitofish, threespine sticklebacks and some bullheads), total length will be measured laterally along the mid-line to the posterior edge of the tail. The weights of fish will consist of measurements to the nearest 0.1 gram. Measure and weigh one fish at a time. Hands, dip nets, and measuring boards should always be wet before coming in contact with fish. Weight measurements should be the final step in the sampling process to allow for expulsion of retained water. If more than 100 fall-run Chinook salmon or more than 50 individuals belonging to a non-salmon species are caught on a given day, then subsampling procedures should be used to acquire data from a subset of the day's catch so there is not a need to devote several hours to data collection activities. The process for conducting subsampling is described in the "Sub-sampling Protocol" section above. For salmon that appear to be spring- or winter-run Chinook salmon based on the length-at-date criteria, the collection of biological samples from those fish may be necessary for exploratory purposes. It is unlikely that these fish are indeed spring- or winter-run Chinook salmon, a fin-clip sample will be used to verify the run that fall outside of the fall-run LAD criteria. The protocol for collecting fin clip samples is described in the "Collection of Fin Clips and Whole Fish" section below.

Any fish present in the tub or trap live well in excess of the above indicated total listed for their species, that are not measured and weighed should be recorded on the appropriate data sheets as a "live plus count" or "mort plus count" for that species. A "plus count" was defined as the total number of fish that were caught in a trap on a given day, and that were not measured, weighed, or assigned a life stage. However, plus counts are still checked for marks and enumerated.

If river temperatures exceed 20°C, enumerate all listed species and release them. Do not hold for fork lengths. On a case by case basis, if temperatures are high and Chinook salmon and steelhead are not staying alive with frequent water changes, air bubblers and frozen water bottles just enumerate them and release. If possible keep a maximum of 50 of each species for fork lengths, if and only if, holding 50 is manageable for the crew to keep alive.

Salmon that are measured for fork length, weight and life stage should be anesthetized prior to measuring using the procedures described in the "Anesthetizing Salmon That Are Measured" section below. If a fin clip is to be taken from a juvenile salmon, that fin clip should be collected after the salmon has been anesthetized, and fork length, weight, and life stage have been recorded.

If a mark-recapture trap efficiency trial is scheduled to occur within the next 48 hours following initial capture, field biologists should consider whether captured salmon should be retained for use in an efficiency trial. If they are not, those salmon can be released below the trap after they have been counted and measured. All other fish species should be released below the RSTs after they have been processed.

Anesthetizing Salmon That Are Measured

Juvenile salmonids that are assessed for fork length, weight, and life stage should always be anesthetized prior to measuring. To anesthetize fish, a solution with Alka-Seltzer is used. In general, fish are immersed in a bath of Alka-Seltzer at a concentration of one tablet for each liter of water. The action of the anesthesia is readily reversed when fish are transferred to fresh water.

The effectiveness and effect of premixed solutions is related to a variety of factors including concentration, fish size, water temperature, stock solution age, and exposure to sunlight. Overexposure (in time or concentration) to the Alka-Seltzer solution will lead to death of fish. Biologists should routinely observe the gill activity of fish immersed in an anesthesia solution; if fish are found to no longer possess, or have markedly reduced gill activity, they should immediately be transferred to fresh water so they can recover.

As fish are measured and are exposed to the solution of Alka-Seltzer, fish should be processed in batches of ~25 individuals to avoid anesthetizing too many fish at one time. Fish size and the crew member's quickness in measuring fish are factors that should be considered when determining how many fish to anesthetize at once.

After fish have been measured they should immediately be placed in a 5 gallon bucket of fresh water for recovery. This bucket should contain 5 teaspoons of Poly Aqua to aid in recovery. Poly Aqua will aid in the regeneration of their slime coat that may have shed during capture and handling. Once fish have fully recovered, release them back in the river.

Collection of Fin Clips

Fin clips will be collected from large juvenile salmon that key out to be spring- or winter-run Chinook salmon according to the length-at-date criteria. Those samples will ultimately be used to assess the salmon run of the individuals the fin clips were taken from. The fin clips should be collected by taking a small pair of surgical scissors and removing not more than $\frac{1}{4}$ of the upper lobe of the caudal fin. The fin clip should then be placed in a vial with 200 proof ethanol, and placed in an envelope with the following information: (1) Species; (2) pre-designated sample number from alcohol vial; (3) sampling location, i.e., Stanislaus River; (4) the putative salmon run designation based on length-at-date criteria; (5) name of the biologist collecting the fin clip; (6) fork length and weight; (7) mortality; (8) life stage; (9) date the sample was collected, and (10) gear (trap and subsite code fish was captured in).

The vials containing fin clips should be stored in a manner that allows for successful retrieval.

Field Quality Check

The first step of data quality assurance/quality check (QA/QC) happens in the field. After all the data have been collected during a sampling visit, each data sheet should be reviewed before the biologists leave the trap site to make sure all information is complete, and any missing values are collected. Common errors include leaving blanks on the data sheet, illegible entries, clarity of plus counts, incorrect species or station codes, and unclear comments. The field quality check should occur before leaving the site so additional data can be collected if necessary. Do not leave data sheets in vehicles or in clipboards as they may get lost or damaged, and return the completed data sheets to the office the same day they are filled out.

Trap Maintenance

Before the field crew leaves the traps at the end of a sample period, they should inspect:

1. The live-box seal for any cracks and proper seating around the trap cone.
2. The trap cone shaft and bushings for cracks and abnormal or excessive wear.
3. The cone's screen for any tears, separation from the cone frame, and the access doors for proper closure. Rivets that attach the screen to the frame often need replacing throughout the season.
4. The winch system, including the cable and pulley, for proper function.
5. The counter system for proper function.
6. The anchor points and cabling system for weaknesses/non-secure attachment.
7. The collars are firmly attached and the collar bolt is tight. The collar should not spin independently of the axle.

MARK/RECAPTURE TRIALS

RSTs only capture a small fraction of the total number of fish migrating past a trap. To estimate the total number of fish produced by the Stanislaus River, it is necessary to conduct mark/recapture trials that quantify the percentage of the total fish population sampled by the traps. These trials should, under ideal circumstances, be conducted in a manner that reduces sampling bias, and are done whenever there are substantial environmental changes that could affect the efficiency of the traps, e.g., a marked increase in stream discharge. The potential source of bias that could affect the accuracy of the trap efficiency tests are as follows:

1. The behavior of the fish that are captured on a daily basis is different than fish used in trap efficiency trials.
2. The trap efficiency test fish are not recognized as they are recaptured.
3. The salmon released during trap efficiency tests do not mix with non-trap efficiency test fish when released.

To conduct the trap efficiency trials fish are marked with a Bismark brown *Y* stain (BBY) and/or a colored elastomer dye. It is therefore imperative that every fish that is captured each day be examined to determine if it has some mark which would indicate it was part of a trap efficiency test.

Under ideal circumstances, wild salmon and not hatchery salmon should be used to conduct trap efficiency tests. Wild salmon that are caught with the RSTs should therefore be used to conduct trap efficiency tests whenever possible. Field staff should attempt to conduct trap efficiency tests on a schedule of once a week, or when scouring events or other events affecting trap efficiency occur. With that sampling in mind, field staff should retain captured wild fish and sequester them in live carts that are held in the RST live-boxes until such time that a sufficient

number of salmon have been collected to conduct a trap efficiency test. Based on historical trap efficiency results, field staff should strive to release at least 1,000 fish during a given efficiency test. Releasing a smaller number of fish has historically resulted in too few recaptures to make the trap efficiency test worthwhile.

To create the ability to discriminate between different groups of marked salmon that were marked as part of a trap efficiency test, it is critical that the marking patterns be applied in a manner and combination where biologists can successfully discriminate between different groups of marked fish. For that reason, the same marking combination (e.g., applying a green mark to the upper caudal fin) should never be used within the same 30-day period when using the colored elastomer dye. Combinations of marking patterns that could cause confusion in proper identification, e.g., applying a green mark to the upper caudal fin one week and then applying a blue mark to the upper caudal fin the following week should also be avoided. It is also important to recognize that some biologists may be color blind, which would therefore compromise their ability to accurately identify the color of the marks applied to fish. For that reason, each biologist should be checked before the start of a field season to determine if they are color blind. If they are, those individuals should not be assigned to processing fish, and they should instead serve in a data recorder capacity when they are in the field.

Typically, fish < 50 mm must be dyed with the BBY whole body stain due to their size. The elastomer dye will typically be difficult to apply to fry due to the translucence and delicate nature of their fins. Apply a whole body stain to small salmon < 50 mm in fork length as described in APPENDIX F: Process for Staining Chinook Salmon with Bismark Brown Dye.

Apply a colored elastomer dye to larger salmon > 50 mm in fork length as described in APPENDIX G: Process for Marking Chinook Salmon with Elastomer Dye.

A few hours prior to the release of salmon used in a trap efficiency test, check each marked fish to determine mark retention and mortality. On the Marked Chinook Salmon Data Sheet, quantify and record the number of salmon that were successfully dyed and stained, and quantify and record the number that were not successfully dyed and stained. Take a random sample of 100 of the successfully dyed and stained salmon and measure and record their fork lengths on the Marked Chinook Salmon Data Sheet.

Additional care must be taken when transporting fish from the hatchery to the release site. The Merced Fish Hatchery (MFH) closes at 3:30 PM, so all of the marking work must be completed by that time. Transportation of fish is done with the use of 3-5 coolers with aeration, with fish being divided evenly between the coolers so as to ensure that fish are not being overcrowded. The appropriate dosage of slime coat protectant should be added to each cooler to ensure fish health. On warmer days, ice may need to be added to fish holding containers to keep the holding water from becoming too warm. Ice can be provided by staff at the MFH if they are asked to do so. The DO and water temperature in the transport containers should be noted before leaving the hatchery. Keep in mind, travel time from the hatchery to the release site is typically 30 minutes, but may be up to 45 minutes to an hour if local traffic is congested. Once at the release site, fish should be transferred from coolers to live carts and set into the river. Again, appropriate amount

of live carts must be used to prevent overcrowding. Once in the live cart and placed in the river, fish are able to sit in the live carts safely for several hours.

On the day when marked salmon are to be released, they should be netted out of coolers and held in live carts full of river water. Each live cart needs to be closely examined for dead, nearly dead, or weak swimming fish. Such fish should be counted and removed from the live marked fish that are to be released. Field staff should make sure they write the number of weak or dead fish in on the Mark and Release datasheet and that number is subtracted from the number of total fish marked to quantify the total number released. Keep the weak and/or dead fish and do not release them in the river where they can be recaptured; they should instead be released downstream of the westernmost trap. Also, record the time fish were released.

One live cart of live, successfully marked salmon should be released on the north half of the river, and the other half released on the south half of the river. Fish should not be “dumped” into the water, rather, they should be released in small net fulls (piecemeal) so that they don’t have an opportunity to “school” as they move downstream. This helps to meet the assumption of random distribution with unmarked fish.

Generally fish are released at the release site at twilight. Therefore, field staff should plan accordingly to ensure they can conduct all the necessary activities to release fish by sunset.

Avoid running the boat between the release site and the RSTs after the fish have been released. If a boat is used to release fish and must travel downstream after a release, remain at the release site for ~15 minutes, then float or row downstream.

At the time of release, make sure the following data are known and recorded on the Mark and Release datasheet: (1) release date and time, (2) number of fish marked, (3) number of mortalities, (4) total number of fish released, and (5) mark type and color.

On mornings following the release of marked salmon and during regular trap checks, the crew should carefully check EVERY fish for a mark. If a mark is observed, that fish must be saved for processing. Marked fish are recorded on data sheets labeled “Recaptured Chinook Salmon Data Form” separate from the “Unmarked Chinook Salmon Data Form.” In addition, each type of marking must be further separated by their marking type or color which signifies the particular fish being a part of a separate release group. Fork length, life stage, race, and mortality for recaptured salmon are recorded on the “Recaptured Chinook Salmon Data Form” for the first 100 random recaptured Chinook salmon; additional recaptured Chinook salmon will be enumerated and tallied under the “Live Unmeasured Plus Count” or “Morts Unmeasured Plus Count” section. Marked Chinook salmon may be observed weeks later, so every fish must be carefully examined even if it’s been many days/weeks since the last marked trial was released.

Make sure the Mark and Release datasheet is completely filled out when staining or releasing Chinook salmon. That datasheet should be delivered to the data crew lead as soon as possible.

COMPUTER DATA ENTRY AND MANAGEMENT

Data sheets need to be delivered to the data crew lead or placed in the “Completed Datasheets” paper tray before the end of each work day.

Data Entry

Data will be stored and analyzed in the Comprehensive Assessment and Monitoring Program’s rotary screw trap platform (Platform). Data should be entered in the platform as soon as possible after collection, ideally on a daily basis. Care should be taken to assure data are entered correctly.

QA/QC Procedure

Ensuring that field data is entered into the Platform so that they are complete and accurate is essential to data management. The Platform and its associated QC/QC database has a number of tools that were developed to look for problems in the quality of the data and the data entry process. Staff that enters data into the Platform should refer to, and become familiar with the user manuals that were prepared for the Platform and the QC/QC database; doing so will provide the ability to detect and then correct problems with the RST data.

First, data sheets will be checked to make sure the data sheets have been completely filled out and there are no errors before data are entered into the CAMP platform. Any corrections made during this process will be made in red ink or pencil. The datasheets are then initialed and dated to indicate they have been checked. Then data will be entered into the CAMP platform by the data crew lead and then QA/QC’d by crew members to ensure that data has been entered correctly.

The verification process will check for entry errors by comparing data sheets with queries that produce hard copy reports from the platform. Corrections, if needed, will be filled out on the QA/QC Data Sheet by crew members and the data crew lead will correct entries in the CAMP Platform. As data is checked, QA/QC data sheets will be signed with initials of the person(s) checking data and the date verified.

Data entry error example: in the course of reviewing the measurements of the randomly selected juvenile salmon that were measured on day X, a salmon is found that is outside the typical size range for a fall-run Chinook salmon on day X based on the length-at-date criteria, the record for that salmon needs to be changed to reflect the appropriate salmon run according to the length-at-date criteria for the day that fish was captured. The person who finds this error should report it on the QA/QC data sheet with their initials, and date verified. Then return the QA/QC data sheet to the data crew lead so that the CAMP database can be updated accordingly.

An effort should be made to verify data on a weekly basis. The QA/QC routines in the platform should be executed and the reports associated with those QA/QC routines should be checked to look for missing data, out of sequence records, and biological attributes that are unusual or abnormal.

After the data has been verified by the crew, the data crew lead will use the QA/QC database provided by Connie Shannon to further check for any additional data entry errors that may have occurred during the data entry process.

APPENDIX A: Field Data Sheets

Trap Visit Data Sheet. Used to record environmental data, trap data, and summarize catch totals.

STANISLAUS RIVER, CASWELL STATE PARK

Field Crew:
 Recording Data: _____
 Measuring Fish: _____

Effort:
 Visit Time: ____/____/____ @ ____:____ AM / PM
Start
 Visit Time 2: ____/____/____ @ ____:____ AM / PM
Stop
 Visit Type ID: _____

Debris:
 Trap Functioning ID: _____
(1: Normal, 2: Functioning but not normal, 3: Stopped functioning, 4: Not in service)
 Total clicker revs if trap stopped: _____

Dominant Debris Type: _____

Debris Volume: _____ (gal)
(Very heavy (60-61), Heavy (40-61), Medium (40-21), Light (20-1), None)

Intakes: ____/____
(0=Completely blocked; backed up into cone, 2=Blocked, 1=Partially blocked, 0=Clear)

Revolutions:
 Before Cleaning RPM: _____ Avg: _____
 After Cleaning RPM: _____ Avg: _____
 Total revs from clicker: _____

Velocity:
 Velocity (nearest 0.01 m/s): _____
(Measured @ 2 ft depth in front of cone on 8 ft trap)

Conditions:
 River Depth: _____ cm Cone Depth: 122 + - _____ cm
 Staff Gage: _____ cm @ ____:____ AM / PM
 River Temp: _____ °C DO: _____ (mg/L)
 Checked Thermograph: _____ (0=No 1=Yes)
 Water sample taken: _____ (0=No 1=Yes)
 NTU: _____ & _____ Avg: _____

Weather: clear partlycloudy overcast foggy
 sprinkle rain heavyrain
 slightlywindy windy
(Circle what best describes the conditions for the day)

Air Temp High: _____ °F and Low: _____ °F
(weather.com)

Notes:

DATE: ____/____/____ Trap ID: _____ Pg ____/____

1. Today's Catch of UNMARKED juvenile Chinook Salmon

Chinook	Live Measured	Live Unmeasured	Morts Measured	Morts Unmeasured	Total
Fall					
Spring					
Winter					
Late fall					
CS Total					

2. Today's Catch of MARKED Chinook Salmon

Chinook	Live Measured	Live Unmeasured	Morts Measured	Morts Unmeasured	Total
CS ad-clipped					
CS BBY recap					
CS Photonic Mark Color					

3. Today's Catch of UNMARKED Steelhead

Steelhead	Live Measured	Live Unmeasured	Morts Measured	Morts Unmeasured	Total
YOY					
Yearling					
Adult					
SH Total					

4. Today's Catch of MARKED Steelhead

Steelhead	Live Measured	Live Unmeasured	Morts Measured	Morts Unmeasured	Total
SH ad-clipped					

Today's Chinook Salmon Race Designation Chart

Race	Minimum	Maximum
Fall		
Spring		
Winter		
Late Fall		

Unmarked Chinook Salmon Data Sheet. Used to record a random sample of 100 Chinook salmon.

Randomly Selected Unmarked Chinook Salmon							DATE: _____	
							TRAP ID: _____ PG: ____ / ____	
Fall-run: (____ - ____)		Spring-run: (____ - ____)		Winter-run: (____ - ____)		Late-fall-run: (____ - ____)		
#	FL	WW	STAGE	Race	MORT	ID #		
1			1 2 3 4 5	F S W L	Y N			
2			1 2 3 4 5	F S W L	Y N			
3			1 2 3 4 5	F S W L	Y N			
4			1 2 3 4 5	F S W L	Y N			
5			1 2 3 4 5	F S W L	Y N			
6			1 2 3 4 5	F S W L	Y N			
7			1 2 3 4 5	F S W L	Y N			
8			1 2 3 4 5	F S W L	Y N			
9			1 2 3 4 5	F S W L	Y N			
10			1 2 3 4 5	F S W L	Y N			
11			1 2 3 4 5	F S W L	Y N			
12			1 2 3 4 5	F S W L	Y N			
13			1 2 3 4 5	F S W L	Y N			
14			1 2 3 4 5	F S W L	Y N			
15			1 2 3 4 5	F S W L	Y N			
16			1 2 3 4 5	F S W L	Y N			
17			1 2 3 4 5	F S W L	Y N			
18			1 2 3 4 5	F S W L	Y N			
19			1 2 3 4 5	F S W L	Y N			
20			1 2 3 4 5	F S W L	Y N			
21			1 2 3 4 5	F S W L	Y N			
22			1 2 3 4 5	F S W L	Y N			
23			1 2 3 4 5	F S W L	Y N			
24			1 2 3 4 5	F S W L	Y N			
25			1 2 3 4 5	F S W L	Y N			
26			1 2 3 4 5	F S W L	Y N			
27			1 2 3 4 5	F S W L	Y N			
28			1 2 3 4 5	F S W L	Y N			
29			1 2 3 4 5	F S W L	Y N			
30			1 2 3 4 5	F S W L	Y N			
31			1 2 3 4 5	F S W L	Y N			
32			1 2 3 4 5	F S W L	Y N			
33			1 2 3 4 5	F S W L	Y N			
34			1 2 3 4 5	F S W L	Y N			
35			1 2 3 4 5	F S W L	Y N			
36			1 2 3 4 5	F S W L	Y N			
37			1 2 3 4 5	F S W L	Y N			
38			1 2 3 4 5	F S W L	Y N			
39			1 2 3 4 5	F S W L	Y N			
40			1 2 3 4 5	F S W L	Y N			
41			1 2 3 4 5	F S W L	Y N			
42			1 2 3 4 5	F S W L	Y N			
43			1 2 3 4 5	F S W L	Y N			
44			1 2 3 4 5	F S W L	Y N			
45			1 2 3 4 5	F S W L	Y N			
46			1 2 3 4 5	F S W L	Y N			
47			1 2 3 4 5	F S W L	Y N			
48			1 2 3 4 5	F S W L	Y N			
49			1 2 3 4 5	F S W L	Y N			
50			1 2 3 4 5	F S W L	Y N			
#	FL	WW	STAGE	Race	MORT	ID #		
51			1 2 3 4 5	F S W L	Y N			
52			1 2 3 4 5	F S W L	Y N			
53			1 2 3 4 5	F S W L	Y N			
54			1 2 3 4 5	F S W L	Y N			
55			1 2 3 4 5	F S W L	Y N			
56			1 2 3 4 5	F S W L	Y N			
57			1 2 3 4 5	F S W L	Y N			
58			1 2 3 4 5	F S W L	Y N			
59			1 2 3 4 5	F S W L	Y N			
60			1 2 3 4 5	F S W L	Y N			
61			1 2 3 4 5	F S W L	Y N			
62			1 2 3 4 5	F S W L	Y N			
63			1 2 3 4 5	F S W L	Y N			
64			1 2 3 4 5	F S W L	Y N			
65			1 2 3 4 5	F S W L	Y N			
66			1 2 3 4 5	F S W L	Y N			
67			1 2 3 4 5	F S W L	Y N			
68			1 2 3 4 5	F S W L	Y N			
69			1 2 3 4 5	F S W L	Y N			
70			1 2 3 4 5	F S W L	Y N			
71			1 2 3 4 5	F S W L	Y N			
72			1 2 3 4 5	F S W L	Y N			
73			1 2 3 4 5	F S W L	Y N			
74			1 2 3 4 5	F S W L	Y N			
75			1 2 3 4 5	F S W L	Y N			
76			1 2 3 4 5	F S W L	Y N			
77			1 2 3 4 5	F S W L	Y N			
78			1 2 3 4 5	F S W L	Y N			
79			1 2 3 4 5	F S W L	Y N			
80			1 2 3 4 5	F S W L	Y N			
81			1 2 3 4 5	F S W L	Y N			
82			1 2 3 4 5	F S W L	Y N			
83			1 2 3 4 5	F S W L	Y N			
84			1 2 3 4 5	F S W L	Y N			
85			1 2 3 4 5	F S W L	Y N			
86			1 2 3 4 5	F S W L	Y N			
87			1 2 3 4 5	F S W L	Y N			
88			1 2 3 4 5	F S W L	Y N			
89			1 2 3 4 5	F S W L	Y N			
90			1 2 3 4 5	F S W L	Y N			
91			1 2 3 4 5	F S W L	Y N			
92			1 2 3 4 5	F S W L	Y N			
93			1 2 3 4 5	F S W L	Y N			
94			1 2 3 4 5	F S W L	Y N			
95			1 2 3 4 5	F S W L	Y N			
96			1 2 3 4 5	F S W L	Y N			
97			1 2 3 4 5	F S W L	Y N			
98			1 2 3 4 5	F S W L	Y N			
99			1 2 3 4 5	F S W L	Y N			
100			1 2 3 4 5	F S W L	Y N			

Unmarked Steelhead Data Sheet. Used to record a random sample of 100 steelhead.

Unmarked Steelhead (Random)

DATE: _____

TRAP ID: _____ Pg: ____ / ____

Steelhead									
#	FL	WW	STAGE						MORT
1			1	2	3	4	5	6	Y N
2			1	2	3	4	5	6	Y N
3			1	2	3	4	5	6	Y N
4			1	2	3	4	5	6	Y N
5			1	2	3	4	5	6	Y N
6			1	2	3	4	5	6	Y N
7			1	2	3	4	5	6	Y N
8			1	2	3	4	5	6	Y N
9			1	2	3	4	5	6	Y N
10			1	2	3	4	5	6	Y N
11			1	2	3	4	5	6	Y N
12			1	2	3	4	5	6	Y N
13			1	2	3	4	5	6	Y N
14			1	2	3	4	5	6	Y N
15			1	2	3	4	5	6	Y N
16			1	2	3	4	5	6	Y N
17			1	2	3	4	5	6	Y N
18			1	2	3	4	5	6	Y N
19			1	2	3	4	5	6	Y N
20			1	2	3	4	5	6	Y N
21			1	2	3	4	5	6	Y N
22			1	2	3	4	5	6	Y N
23			1	2	3	4	5	6	Y N
24			1	2	3	4	5	6	Y N
25			1	2	3	4	5	6	Y N
26			1	2	3	4	5	6	Y N
27			1	2	3	4	5	6	Y N
28			1	2	3	4	5	6	Y N
29			1	2	3	4	5	6	Y N
30			1	2	3	4	5	6	Y N
31			1	2	3	4	5	6	Y N
32			1	2	3	4	5	6	Y N
33			1	2	3	4	5	6	Y N
34			1	2	3	4	5	6	Y N
35			1	2	3	4	5	6	Y N
36			1	2	3	4	5	6	Y N
37			1	2	3	4	5	6	Y N
38			1	2	3	4	5	6	Y N
39			1	2	3	4	5	6	Y N
40			1	2	3	4	5	6	Y N
41			1	2	3	4	5	6	Y N
42			1	2	3	4	5	6	Y N
43			1	2	3	4	5	6	Y N
44			1	2	3	4	5	6	Y N
45			1	2	3	4	5	6	Y N
46			1	2	3	4	5	6	Y N
47			1	2	3	4	5	6	Y N
48			1	2	3	4	5	6	Y N
49			1	2	3	4	5	6	Y N
50			1	2	3	4	5	6	Y N

Live Unmeasured Plus Count

Total Live Unmeasured

Steelhead									
#	FL	WW	STAGE						MORT
51			1	2	3	4	5	6	Y N
52			1	2	3	4	5	6	Y N
53			1	2	3	4	5	6	Y N
54			1	2	3	4	5	6	Y N
55			1	2	3	4	5	6	Y N
56			1	2	3	4	5	6	Y N
57			1	2	3	4	5	6	Y N
58			1	2	3	4	5	6	Y N
59			1	2	3	4	5	6	Y N
60			1	2	3	4	5	6	Y N
61			1	2	3	4	5	6	Y N
62			1	2	3	4	5	6	Y N
63			1	2	3	4	5	6	Y N
64			1	2	3	4	5	6	Y N
65			1	2	3	4	5	6	Y N
66			1	2	3	4	5	6	Y N
67			1	2	3	4	5	6	Y N
68			1	2	3	4	5	6	Y N
69			1	2	3	4	5	6	Y N
70			1	2	3	4	5	6	Y N
71			1	2	3	4	5	6	Y N
72			1	2	3	4	5	6	Y N
73			1	2	3	4	5	6	Y N
74			1	2	3	4	5	6	Y N
75			1	2	3	4	5	6	Y N
76			1	2	3	4	5	6	Y N
77			1	2	3	4	5	6	Y N
78			1	2	3	4	5	6	Y N
79			1	2	3	4	5	6	Y N
80			1	2	3	4	5	6	Y N
81			1	2	3	4	5	6	Y N
82			1	2	3	4	5	6	Y N
83			1	2	3	4	5	6	Y N
84			1	2	3	4	5	6	Y N
85			1	2	3	4	5	6	Y N
86			1	2	3	4	5	6	Y N
87			1	2	3	4	5	6	Y N
88			1	2	3	4	5	6	Y N
89			1	2	3	4	5	6	Y N
90			1	2	3	4	5	6	Y N
91			1	2	3	4	5	6	Y N
92			1	2	3	4	5	6	Y N
93			1	2	3	4	5	6	Y N
94			1	2	3	4	5	6	Y N
95			1	2	3	4	5	6	Y N
96			1	2	3	4	5	6	Y N
97			1	2	3	4	5	6	Y N
98			1	2	3	4	5	6	Y N
99			1	2	3	4	5	6	Y N
100			1	2	3	4	5	6	Y N

Morts Unmeasured Plus Count

Total Morts Unmeasured

Spring-run and Winter-run LAD Chinook Salmon Fin Clip Data Sheet. Used to record data on spring-run and winter-run (by LAD) Chinook salmon when fin clip samples are taken.

Spring/Winter Chinook Salmon Upper Caudal Fin Clips (Not Random) DATE: _____					
U.M.= Unmeasured					
TRAP ID: _____ PG: ____/____					
RACE		Upper Caudal Fin Clips			
#	FL	WW	STAGE	MORT	ID#
1			1 2 3 4 5	Y N	
2			1 2 3 4 5	Y N	
3			1 2 3 4 5	Y N	
4			1 2 3 4 5	Y N	
5			1 2 3 4 5	Y N	
6			1 2 3 4 5	Y N	
7			1 2 3 4 5	Y N	
8			1 2 3 4 5	Y N	
9			1 2 3 4 5	Y N	
10			1 2 3 4 5	Y N	
11			1 2 3 4 5	Y N	
12			1 2 3 4 5	Y N	
13			1 2 3 4 5	Y N	
14			1 2 3 4 5	Y N	
15			1 2 3 4 5	Y N	
16			1 2 3 4 5	Y N	
17			1 2 3 4 5	Y N	
18			1 2 3 4 5	Y N	
19			1 2 3 4 5	Y N	
20			1 2 3 4 5	Y N	
21			1 2 3 4 5	Y N	
22			1 2 3 4 5	Y N	
23			1 2 3 4 5	Y N	
24			1 2 3 4 5	Y N	
25			1 2 3 4 5	Y N	
26			1 2 3 4 5	Y N	
27			1 2 3 4 5	Y N	
28			1 2 3 4 5	Y N	
29			1 2 3 4 5	Y N	
30			1 2 3 4 5	Y N	
31			1 2 3 4 5	Y N	
32			1 2 3 4 5	Y N	
33			1 2 3 4 5	Y N	
34			1 2 3 4 5	Y N	
35			1 2 3 4 5	Y N	
36			1 2 3 4 5	Y N	
37			1 2 3 4 5	Y N	
38			1 2 3 4 5	Y N	
39			1 2 3 4 5	Y N	
40			1 2 3 4 5	Y N	
41			1 2 3 4 5	Y N	
42			1 2 3 4 5	Y N	
43			1 2 3 4 5	Y N	
44			1 2 3 4 5	Y N	
45			1 2 3 4 5	Y N	
46			1 2 3 4 5	Y N	
47			1 2 3 4 5	Y N	
48			1 2 3 4 5	Y N	
49			1 2 3 4 5	Y N	
50			1 2 3 4 5	Y N	
Live U.M. Plus Count				Morts U.M. Plus Count	
Total Live U.M.=				Total Morts U.M.=	

RACE		Upper Caudal Fin Clips			
#	FL	WW	STAGE	MORT	ID#
1			1 2 3 4 5	Y N	
2			1 2 3 4 5	Y N	
3			1 2 3 4 5	Y N	
4			1 2 3 4 5	Y N	
5			1 2 3 4 5	Y N	
6			1 2 3 4 5	Y N	
7			1 2 3 4 5	Y N	
8			1 2 3 4 5	Y N	
9			1 2 3 4 5	Y N	
10			1 2 3 4 5	Y N	
11			1 2 3 4 5	Y N	
12			1 2 3 4 5	Y N	
13			1 2 3 4 5	Y N	
14			1 2 3 4 5	Y N	
15			1 2 3 4 5	Y N	
16			1 2 3 4 5	Y N	
17			1 2 3 4 5	Y N	
18			1 2 3 4 5	Y N	
19			1 2 3 4 5	Y N	
20			1 2 3 4 5	Y N	
21			1 2 3 4 5	Y N	
22			1 2 3 4 5	Y N	
23			1 2 3 4 5	Y N	
24			1 2 3 4 5	Y N	
25			1 2 3 4 5	Y N	
26			1 2 3 4 5	Y N	
27			1 2 3 4 5	Y N	
28			1 2 3 4 5	Y N	
29			1 2 3 4 5	Y N	
30			1 2 3 4 5	Y N	
31			1 2 3 4 5	Y N	
32			1 2 3 4 5	Y N	
33			1 2 3 4 5	Y N	
34			1 2 3 4 5	Y N	
35			1 2 3 4 5	Y N	
36			1 2 3 4 5	Y N	
37			1 2 3 4 5	Y N	
38			1 2 3 4 5	Y N	
39			1 2 3 4 5	Y N	
40			1 2 3 4 5	Y N	
41			1 2 3 4 5	Y N	
42			1 2 3 4 5	Y N	
43			1 2 3 4 5	Y N	
44			1 2 3 4 5	Y N	
45			1 2 3 4 5	Y N	
46			1 2 3 4 5	Y N	
47			1 2 3 4 5	Y N	
48			1 2 3 4 5	Y N	
49			1 2 3 4 5	Y N	
50			1 2 3 4 5	Y N	
Live U.M. Plus Count				Morts U.M. Plus Count	
Total Live U.M.=				Total Morts U.M.=	

Photonically Marked Chinook Salmon Data Sheet. Used to record data on hatchery Chinook salmon that have been marked with photonic fluorescent dye used for a trap efficiency trial.

Merced Hatchery Fall-run Chinook Salmon
Photonic Marking Data Sheet
 Photonic Color: Photonic Green=PG Photonic Pink=PP Photonic Orange=PO

DATE: ____/____/____

#	FL	STAGE	MORT
1		1 2 3 4 5	Y N
2		1 2 3 4 5	Y N
3		1 2 3 4 5	Y N
4		1 2 3 4 5	Y N
5		1 2 3 4 5	Y N
6		1 2 3 4 5	Y N
7		1 2 3 4 5	Y N
8		1 2 3 4 5	Y N
9		1 2 3 4 5	Y N
10		1 2 3 4 5	Y N
11		1 2 3 4 5	Y N
12		1 2 3 4 5	Y N
13		1 2 3 4 5	Y N
14		1 2 3 4 5	Y N
15		1 2 3 4 5	Y N
16		1 2 3 4 5	Y N
17		1 2 3 4 5	Y N
18		1 2 3 4 5	Y N
19		1 2 3 4 5	Y N
20		1 2 3 4 5	Y N
21		1 2 3 4 5	Y N
22		1 2 3 4 5	Y N
23		1 2 3 4 5	Y N
24		1 2 3 4 5	Y N
25		1 2 3 4 5	Y N
26		1 2 3 4 5	Y N
27		1 2 3 4 5	Y N
28		1 2 3 4 5	Y N
29		1 2 3 4 5	Y N
30		1 2 3 4 5	Y N
31		1 2 3 4 5	Y N
32		1 2 3 4 5	Y N
33		1 2 3 4 5	Y N
34		1 2 3 4 5	Y N
35		1 2 3 4 5	Y N
36		1 2 3 4 5	Y N
37		1 2 3 4 5	Y N
38		1 2 3 4 5	Y N
39		1 2 3 4 5	Y N
40		1 2 3 4 5	Y N
41		1 2 3 4 5	Y N
42		1 2 3 4 5	Y N
43		1 2 3 4 5	Y N
44		1 2 3 4 5	Y N
45		1 2 3 4 5	Y N
46		1 2 3 4 5	Y N
47		1 2 3 4 5	Y N
48		1 2 3 4 5	Y N
49		1 2 3 4 5	Y N
50		1 2 3 4 5	Y N

Live Unmeasured Plus Count

Total Live Unmeasured

#	FL	STAGE	MORT
51		1 2 3 4 5	Y N
52		1 2 3 4 5	Y N
53		1 2 3 4 5	Y N
54		1 2 3 4 5	Y N
55		1 2 3 4 5	Y N
56		1 2 3 4 5	Y N
57		1 2 3 4 5	Y N
58		1 2 3 4 5	Y N
59		1 2 3 4 5	Y N
60		1 2 3 4 5	Y N
61		1 2 3 4 5	Y N
62		1 2 3 4 5	Y N
63		1 2 3 4 5	Y N
64		1 2 3 4 5	Y N
65		1 2 3 4 5	Y N
66		1 2 3 4 5	Y N
67		1 2 3 4 5	Y N
68		1 2 3 4 5	Y N
69		1 2 3 4 5	Y N
70		1 2 3 4 5	Y N
71		1 2 3 4 5	Y N
72		1 2 3 4 5	Y N
73		1 2 3 4 5	Y N
74		1 2 3 4 5	Y N
75		1 2 3 4 5	Y N
76		1 2 3 4 5	Y N
77		1 2 3 4 5	Y N
78		1 2 3 4 5	Y N
79		1 2 3 4 5	Y N
80		1 2 3 4 5	Y N
81		1 2 3 4 5	Y N
82		1 2 3 4 5	Y N
83		1 2 3 4 5	Y N
84		1 2 3 4 5	Y N
85		1 2 3 4 5	Y N
86		1 2 3 4 5	Y N
87		1 2 3 4 5	Y N
88		1 2 3 4 5	Y N
89		1 2 3 4 5	Y N
90		1 2 3 4 5	Y N
91		1 2 3 4 5	Y N
92		1 2 3 4 5	Y N
93		1 2 3 4 5	Y N
94		1 2 3 4 5	Y N
95		1 2 3 4 5	Y N
96		1 2 3 4 5	Y N
97		1 2 3 4 5	Y N
98		1 2 3 4 5	Y N
99		1 2 3 4 5	Y N
100		1 2 3 4 5	Y N

Morts Unmeasured Plus Count

Total Morts Unmeasured

Recaptured Chinook Salmon Data Sheet. Used to record data on Chinook salmon after they've been recaptured from a trap efficiency trial.

Recaptured Chinook Salmon (Not Random)

BBY = Bismark Brown Y PG/PO/PP = Photonic (Color)

U.M.= Unmeasured

DATE: _____

TRAP ID: _____ PG: ____/____

MARK TYPE:				
#	FL	STAGE	MORT	
1		1 2 3 4 5	Y	N
2		1 2 3 4 5	Y	N
3		1 2 3 4 5	Y	N
4		1 2 3 4 5	Y	N
5		1 2 3 4 5	Y	N
6		1 2 3 4 5	Y	N
7		1 2 3 4 5	Y	N
8		1 2 3 4 5	Y	N
9		1 2 3 4 5	Y	N
10		1 2 3 4 5	Y	N
11		1 2 3 4 5	Y	N
12		1 2 3 4 5	Y	N
13		1 2 3 4 5	Y	N
14		1 2 3 4 5	Y	N
15		1 2 3 4 5	Y	N
16		1 2 3 4 5	Y	N
17		1 2 3 4 5	Y	N
18		1 2 3 4 5	Y	N
19		1 2 3 4 5	Y	N
20		1 2 3 4 5	Y	N
21		1 2 3 4 5	Y	N
22		1 2 3 4 5	Y	N
23		1 2 3 4 5	Y	N
24		1 2 3 4 5	Y	N
25		1 2 3 4 5	Y	N
26		1 2 3 4 5	Y	N
27		1 2 3 4 5	Y	N
28		1 2 3 4 5	Y	N
29		1 2 3 4 5	Y	N
30		1 2 3 4 5	Y	N
31		1 2 3 4 5	Y	N
32		1 2 3 4 5	Y	N
33		1 2 3 4 5	Y	N
34		1 2 3 4 5	Y	N
35		1 2 3 4 5	Y	N
36		1 2 3 4 5	Y	N
37		1 2 3 4 5	Y	N
38		1 2 3 4 5	Y	N
39		1 2 3 4 5	Y	N
40		1 2 3 4 5	Y	N
41		1 2 3 4 5	Y	N
42		1 2 3 4 5	Y	N
43		1 2 3 4 5	Y	N
44		1 2 3 4 5	Y	N
45		1 2 3 4 5	Y	N
46		1 2 3 4 5	Y	N
47		1 2 3 4 5	Y	N
48		1 2 3 4 5	Y	N
49		1 2 3 4 5	Y	N
50		1 2 3 4 5	Y	N
Live U.M. Plus Count			Morts U.M. Plus Count	
Total Live U.M.=			Total Morts U.M.=	

MARK TYPE:				
#	FL	STAGE	MORT	
1		1 2 3 4 5	Y	N
2		1 2 3 4 5	Y	N
3		1 2 3 4 5	Y	N
4		1 2 3 4 5	Y	N
5		1 2 3 4 5	Y	N
6		1 2 3 4 5	Y	N
7		1 2 3 4 5	Y	N
8		1 2 3 4 5	Y	N
9		1 2 3 4 5	Y	N
10		1 2 3 4 5	Y	N
11		1 2 3 4 5	Y	N
12		1 2 3 4 5	Y	N
13		1 2 3 4 5	Y	N
14		1 2 3 4 5	Y	N
15		1 2 3 4 5	Y	N
16		1 2 3 4 5	Y	N
17		1 2 3 4 5	Y	N
18		1 2 3 4 5	Y	N
19		1 2 3 4 5	Y	N
20		1 2 3 4 5	Y	N
21		1 2 3 4 5	Y	N
22		1 2 3 4 5	Y	N
23		1 2 3 4 5	Y	N
24		1 2 3 4 5	Y	N
25		1 2 3 4 5	Y	N
26		1 2 3 4 5	Y	N
27		1 2 3 4 5	Y	N
28		1 2 3 4 5	Y	N
29		1 2 3 4 5	Y	N
30		1 2 3 4 5	Y	N
31		1 2 3 4 5	Y	N
32		1 2 3 4 5	Y	N
33		1 2 3 4 5	Y	N
34		1 2 3 4 5	Y	N
35		1 2 3 4 5	Y	N
36		1 2 3 4 5	Y	N
37		1 2 3 4 5	Y	N
38		1 2 3 4 5	Y	N
39		1 2 3 4 5	Y	N
40		1 2 3 4 5	Y	N
41		1 2 3 4 5	Y	N
42		1 2 3 4 5	Y	N
43		1 2 3 4 5	Y	N
44		1 2 3 4 5	Y	N
45		1 2 3 4 5	Y	N
46		1 2 3 4 5	Y	N
47		1 2 3 4 5	Y	N
48		1 2 3 4 5	Y	N
49		1 2 3 4 5	Y	N
50		1 2 3 4 5	Y	N
Live U.M. Plus Count			Morts U.M. Plus Count	
Total Live U.M.=			Total Morts U.M.=	

Adipose Fin Clipped Salmonid Data Sheet. Used to record data on ad-clipped (hatchery origin) Chinook salmon or steelhead.

Adipose Fin Clipped Salmonid (Not Random)

U.M. = Umm easured

DATE: _____

TRAP ID: _____ PG: ____/____

Spp		Ad-Clipped					
#	FL	WW	STAGE	RACE	MORT	ID#	
1			1 2 3 4 5	F S W L Y N			
2			1 2 3 4 5	F S W L Y N			
3			1 2 3 4 5	F S W L Y N			
4			1 2 3 4 5	F S W L Y N			
5			1 2 3 4 5	F S W L Y N			
6			1 2 3 4 5	F S W L Y N			
7			1 2 3 4 5	F S W L Y N			
8			1 2 3 4 5	F S W L Y N			
9			1 2 3 4 5	F S W L Y N			
10			1 2 3 4 5	F S W L Y N			
11			1 2 3 4 5	F S W L Y N			
12			1 2 3 4 5	F S W L Y N			
13			1 2 3 4 5	F S W L Y N			
14			1 2 3 4 5	F S W L Y N			
15			1 2 3 4 5	F S W L Y N			
16			1 2 3 4 5	F S W L Y N			
17			1 2 3 4 5	F S W L Y N			
18			1 2 3 4 5	F S W L Y N			
19			1 2 3 4 5	F S W L Y N			
20			1 2 3 4 5	F S W L Y N			
21			1 2 3 4 5	F S W L Y N			
22			1 2 3 4 5	F S W L Y N			
23			1 2 3 4 5	F S W L Y N			
24			1 2 3 4 5	F S W L Y N			
25			1 2 3 4 5	F S W L Y N			
26			1 2 3 4 5	F S W L Y N			
27			1 2 3 4 5	F S W L Y N			
28			1 2 3 4 5	F S W L Y N			
29			1 2 3 4 5	F S W L Y N			
30			1 2 3 4 5	F S W L Y N			
31			1 2 3 4 5	F S W L Y N			
32			1 2 3 4 5	F S W L Y N			
33			1 2 3 4 5	F S W L Y N			
34			1 2 3 4 5	F S W L Y N			
35			1 2 3 4 5	F S W L Y N			
36			1 2 3 4 5	F S W L Y N			
37			1 2 3 4 5	F S W L Y N			
38			1 2 3 4 5	F S W L Y N			
39			1 2 3 4 5	F S W L Y N			
40			1 2 3 4 5	F S W L Y N			
41			1 2 3 4 5	F S W L Y N			
42			1 2 3 4 5	F S W L Y N			
43			1 2 3 4 5	F S W L Y N			
44			1 2 3 4 5	F S W L Y N			
45			1 2 3 4 5	F S W L Y N			
46			1 2 3 4 5	F S W L Y N			
47			1 2 3 4 5	F S W L Y N			
48			1 2 3 4 5	F S W L Y N			
49			1 2 3 4 5	F S W L Y N			
50			1 2 3 4 5	F S W L Y N			
Live U.M. Plus Count			Morts U.M. Plus Count				
Total Live U.M.=			Total Morts U.M.=				

Spp		Ad-Clipped					
#	FL	WW	STAGE	RACE	MORT	ID#	
1			1 2 3 4 5	F S W L Y N			
2			1 2 3 4 5	F S W L Y N			
3			1 2 3 4 5	F S W L Y N			
4			1 2 3 4 5	F S W L Y N			
5			1 2 3 4 5	F S W L Y N			
6			1 2 3 4 5	F S W L Y N			
7			1 2 3 4 5	F S W L Y N			
8			1 2 3 4 5	F S W L Y N			
9			1 2 3 4 5	F S W L Y N			
10			1 2 3 4 5	F S W L Y N			
11			1 2 3 4 5	F S W L Y N			
12			1 2 3 4 5	F S W L Y N			
13			1 2 3 4 5	F S W L Y N			
14			1 2 3 4 5	F S W L Y N			
15			1 2 3 4 5	F S W L Y N			
16			1 2 3 4 5	F S W L Y N			
17			1 2 3 4 5	F S W L Y N			
18			1 2 3 4 5	F S W L Y N			
19			1 2 3 4 5	F S W L Y N			
20			1 2 3 4 5	F S W L Y N			
21			1 2 3 4 5	F S W L Y N			
22			1 2 3 4 5	F S W L Y N			
23			1 2 3 4 5	F S W L Y N			
24			1 2 3 4 5	F S W L Y N			
25			1 2 3 4 5	F S W L Y N			
26			1 2 3 4 5	F S W L Y N			
27			1 2 3 4 5	F S W L Y N			
28			1 2 3 4 5	F S W L Y N			
29			1 2 3 4 5	F S W L Y N			
30			1 2 3 4 5	F S W L Y N			
31			1 2 3 4 5	F S W L Y N			
32			1 2 3 4 5	F S W L Y N			
33			1 2 3 4 5	F S W L Y N			
34			1 2 3 4 5	F S W L Y N			
35			1 2 3 4 5	F S W L Y N			
36			1 2 3 4 5	F S W L Y N			
37			1 2 3 4 5	F S W L Y N			
38			1 2 3 4 5	F S W L Y N			
39			1 2 3 4 5	F S W L Y N			
40			1 2 3 4 5	F S W L Y N			
41			1 2 3 4 5	F S W L Y N			
42			1 2 3 4 5	F S W L Y N			
43			1 2 3 4 5	F S W L Y N			
44			1 2 3 4 5	F S W L Y N			
45			1 2 3 4 5	F S W L Y N			
46			1 2 3 4 5	F S W L Y N			
47			1 2 3 4 5	F S W L Y N			
48			1 2 3 4 5	F S W L Y N			
49			1 2 3 4 5	F S W L Y N			
50			1 2 3 4 5	F S W L Y N			
Live U.M. Plus Count			Morts U.M. Plus Count				
Total Live U.M.=			Total Morts U.M.=				

Other Species Data Sheet. Used to record data on bycatch (species of fish other than our target taxa).

Other Species
U.M. = Unmeasured

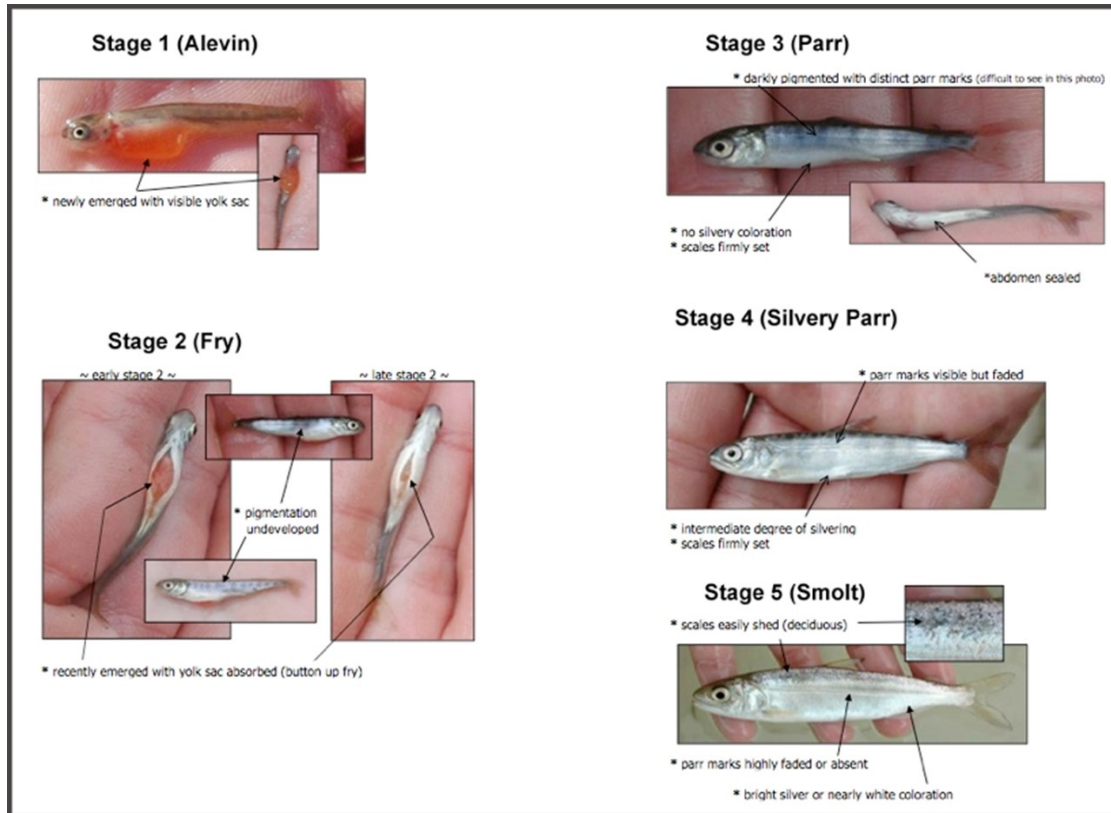
DATE: _____
TRAP ID: _____ PG: ____/____

SPP				SPP				SPP				SPP			
#	FL	STAGE	MORT	#	FL	STAGE	MORT	#	FL	STAGE	MORT	#	FL	STAGE	MORT
1		A J	Y N	1		A J	Y N	1		A J	Y N	1		A J	Y N
2		A J	Y N	2		A J	Y N	2		A J	Y N	2		A J	Y N
3		A J	Y N	3		A J	Y N	3		A J	Y N	3		A J	Y N
4		A J	Y N	4		A J	Y N	4		A J	Y N	4		A J	Y N
5		A J	Y N	5		A J	Y N	5		A J	Y N	5		A J	Y N
6		A J	Y N	6		A J	Y N	6		A J	Y N	6		A J	Y N
7		A J	Y N	7		A J	Y N	7		A J	Y N	7		A J	Y N
8		A J	Y N	8		A J	Y N	8		A J	Y N	8		A J	Y N
9		A J	Y N	9		A J	Y N	9		A J	Y N	9		A J	Y N
10		A J	Y N	10		A J	Y N	10		A J	Y N	10		A J	Y N
11		A J	Y N	11		A J	Y N	11		A J	Y N	11		A J	Y N
12		A J	Y N	12		A J	Y N	12		A J	Y N	12		A J	Y N
13		A J	Y N	13		A J	Y N	13		A J	Y N	13		A J	Y N
14		A J	Y N	14		A J	Y N	14		A J	Y N	14		A J	Y N
15		A J	Y N	15		A J	Y N	15		A J	Y N	15		A J	Y N
16		A J	Y N	16		A J	Y N	16		A J	Y N	16		A J	Y N
17		A J	Y N	17		A J	Y N	17		A J	Y N	17		A J	Y N
18		A J	Y N	18		A J	Y N	18		A J	Y N	18		A J	Y N
19		A J	Y N	19		A J	Y N	19		A J	Y N	19		A J	Y N
20		A J	Y N	20		A J	Y N	20		A J	Y N	20		A J	Y N
21		A J	Y N	21		A J	Y N	21		A J	Y N	21		A J	Y N
22		A J	Y N	22		A J	Y N	22		A J	Y N	22		A J	Y N
23		A J	Y N	23		A J	Y N	23		A J	Y N	23		A J	Y N
24		A J	Y N	24		A J	Y N	24		A J	Y N	24		A J	Y N
25		A J	Y N	25		A J	Y N	25		A J	Y N	25		A J	Y N
26		A J	Y N	26		A J	Y N	26		A J	Y N	26		A J	Y N
27		A J	Y N	27		A J	Y N	27		A J	Y N	27		A J	Y N
28		A J	Y N	28		A J	Y N	28		A J	Y N	28		A J	Y N
29		A J	Y N	29		A J	Y N	29		A J	Y N	29		A J	Y N
30		A J	Y N	30		A J	Y N	30		A J	Y N	30		A J	Y N
31		A J	Y N	31		A J	Y N	31		A J	Y N	31		A J	Y N
32		A J	Y N	32		A J	Y N	32		A J	Y N	32		A J	Y N
33		A J	Y N	33		A J	Y N	33		A J	Y N	33		A J	Y N
34		A J	Y N	34		A J	Y N	34		A J	Y N	34		A J	Y N
35		A J	Y N	35		A J	Y N	35		A J	Y N	35		A J	Y N
36		A J	Y N	36		A J	Y N	36		A J	Y N	36		A J	Y N
37		A J	Y N	37		A J	Y N	37		A J	Y N	37		A J	Y N
38		A J	Y N	38		A J	Y N	38		A J	Y N	38		A J	Y N
39		A J	Y N	39		A J	Y N	39		A J	Y N	39		A J	Y N
40		A J	Y N	40		A J	Y N	40		A J	Y N	40		A J	Y N
41		A J	Y N	41		A J	Y N	41		A J	Y N	41		A J	Y N
42		A J	Y N	42		A J	Y N	42		A J	Y N	42		A J	Y N
43		A J	Y N	43		A J	Y N	43		A J	Y N	43		A J	Y N
44		A J	Y N	44		A J	Y N	44		A J	Y N	44		A J	Y N
45		A J	Y N	45		A J	Y N	45		A J	Y N	45		A J	Y N
46		A J	Y N	46		A J	Y N	46		A J	Y N	46		A J	Y N
47		A J	Y N	47		A J	Y N	47		A J	Y N	47		A J	Y N
48		A J	Y N	48		A J	Y N	48		A J	Y N	48		A J	Y N
49		A J	Y N	49		A J	Y N	49		A J	Y N	49		A J	Y N
50		A J	Y N	50		A J	Y N	50		A J	Y N	50		A J	Y N
Live U.M.		Morts U.M.		Live U.M.		Morts U.M.		Live U.M.		Morts U.M.		Live U.M.		Morts U.M.	
Totals=		Totals=		Totals=		Totals=		Totals=		Totals=		Totals=		Totals=	

APPENDIX B: Chinook Salmon and Steelhead Life Stages

Smolt Index	Life Stage	Criteria
1	Yolk-sac Fry	* Newly emerged with visible yolk sac
2	Fry	* Recently emerged with yolk sac absorbed (button-up fry) * Seam along mid-ventral line visible * Pigmentation undeveloped
3	Parr	* Seam along mid-ventral line not visible * Scales firmly set * Darkly pigmented with distinct parr marks * No silvery coloration
4	Silvery Parr	* Parr marks visible but faded * Intermediate degree of silverying
5	Smolt	* Parr marks highly faded or absent * Bright silver or nearly white coloration * Scales easily shed (deciduous) * Black trailing edge on caudal fin * Body/head elongating
6	Adult	* $\geq 300\text{mm}$

Figures Illustrating Different Chinook Salmon Life Stages



APPENDIX C: Trap Visit Data Sheet Terminology

This appendix defines how particular terms on the field data sheets are defined, and how they should be viewed as data are entered on the field data sheets.

Before cleaning (RPM): the number of revolutions the cone makes in one complete minute, as measured *before* the trap is cleaned. Take three separate RPM readings and then calculate and record the average. The before revs do not need to be recorded if the trap is stopped on arrival.

The determination of when a trap makes a complete revolution is made by observing a specific location on the trap, e.g., a colored dot or bolt present on the trap cone. On most 8-foot RST trap cones, there is one weld line on the base of cone and 10 equally sized screen sections which makes it simple to reference a full rotation and 1/10 of a rotation.

After cleaning (RPM): the number of revolutions the cone makes in one complete minute, as measured *after* the trap is cleaned. Take three separate RPM readings and then calculate and record the average.

Cone Depth (inches): this parameter needs to be quantified before staff board the RST. After personnel have boarded the trap, the cone depth may change as much as three inches. It's therefore crucial to take this reading before the boat is secured and personnel board the trap. The cone depth gauge is located directly forward of the cone and mounted on the inside of the pontoon. The cone depth gauge has one inch increments ranging from +4 inches to -4 inches. Your measurement will either be added to or subtracted from the standard cone depth of 48 inches (the radius of an eight foot diameter cone). For example, if the gauge reads minus two inches (-2), then you will record a value of 46 inches on the data sheet for "Cone depth (in)". This parameter is measured at every trap checking/servicing and is never assumed to be the same as it was the day before. The cone depth is used to calculate relative abundance or catch per unit volume (CPUV). A one inch difference makes a change in the number of acre-feet sampled during a sample and, therefore, can have a large impact on CPUV.

Debris Volume (gallons): a description of the total amount of debris in the trap live-box. The amount of debris should be measured using a 5-gallon bucket, and the debris volume on the Trap Visit data sheet should be recorded as the total number of gallons of debris present.

Recorder/crew: initials of the names of the personnel operating the trap. The first set of initials is the data recorder during the trap visit. The subsequent initials represent the individuals that were collecting biological or environmental data.

Sample Gear ID (feet): reflects the cone diameter of a RST at a subsite.

Sample Gear ID	Description
8	if the trap is a 8-foot diameter RST

Subsite code: a code describing the location of a position within the sampling area at Caswell State Park.

River channel	CAMP Subsite code	Trap Visit Data Sheet Subsite Code	Notes
Main	STAN1	8.1	South bank side
Main	STAN2	8.2	North bank side



Total revolutions since last trap check: a lever actuated mechanical counter is mounted on each RST, and records the number of revolutions the trap has been made since the last time the counter was set to a value of 0. The total revolutions since last trap check reflects the reading on the rotation counter at the end of the trap visit before it is reset for the next sampling period. The counter is set to 0 when the crew has determined the live well has been completely cleared of all fish and debris.

Trap Functioning ID: code for how well the trap was functioning when the visit to the trap began. Answer the question, “Did the trap function correctly since the last visit to the trap?”

Trap Functioning ID	Description
1	Trap functioning normally
2	Trap functioning, but not normally (trap has some debris or other impediment that may affect its ability to collect fish, or the trap is not rotating properly)
3	Trap stopped functioning (trap is packed with debris so it can not collect fish, or the trap is not rotating)
4	Trap not in service

Trap Visit Date: the date when the visit to a trap occurred. E.g., if the field crew arrived at a trap on January 1, 2010 to service a trap, the Trap Visit Date would be 01/01/2010.

Visit Time: represents the start of a sample period. The time entered on the data sheet reflects the moment a trap begins fishing.

Visit Time2: represents the time a trap is emptied and is put back in service after it has been cleared of fish. The VisitTime2 field therefore represents the end of a sample period. In most cases when a trap fishes without problems and there is no break in sampling, the VisitTime2 of one day is also the VisitTime for the next day.

For example and when there is no break in sampling, if biologists clear the live box of fish at 1:00 PM on Tuesday, that 1:00 PM Tuesday time is the VisitTime2 data entry for Tuesday’s catch, and it is also the time for the VisitTime of the Wednesday catch period.

An example when there is discontinuous sampling is as follows. If biologists clear a live box of fish at 1:00 PM on Tuesday, and then do not resume trapping until 5:00 PM on Tuesday, then the 1:00 PM time on Tuesday is the VisitTime2 data entry representing the end of the sampling period on Tuesday, and the 5:00 PM time on Tuesday represents the VisitTime of the Wednesday catch period.

Visit Type ID: code for the work that was done during a trap visit. This field is used to help characterize when a trap is started, restarted in the same position and configuration after a malfunction, adjusted, and stopped. Using this field, it is possible to reconstruct the operational history of the trapping at a subsite.

Visit Type ID	Description
1	Start trap and begin trapping. Used when trap had not been operating, or when it is moved or reconfigured. Defines the beginning of a sampling period. Fish are never processed during this type of visit.
2	Continue trapping in same position and configuration without interruption. Used when there is no break in trap operations. Defines the break between two sampling periods. Fish are usually processed during this type of visit.
3	Unplanned restart of trap after malfunction (in same position and configuration). Used when the trap had stopped operating. Defines the break between two sampling periods. Fish are usually processed during this type of visit.
4	End trapping in current position or configuration. Used when trap is moved or stopped. Defines the end of a sampling period. Fish are usually processed during this type of visit.
5	"Drive by", i.e., trap is scanned to ensure it is functioning, but fish are not processed and trap is not adjusted. Environmental measures may be taken. If a "drive by" results in fish being sampled or trap being serviced, then use an alternative code as appropriate. Does not define the beginning or end of a sampling period.
6	Service / adjust / clean trap. Adjustment is made to trap during a sampling period, such as returning it to desired sampling position or removing debris. Does not define the beginning or end of a sampling period. Fish are not processed during this type of visit.

Water dissolved oxygen (milligrams per liter): dissolved oxygen levels in the Stanislaus River will be measured with a YSI® 55 Dissolved Oxygen Meter. The location where dissolved oxygen is measured will be at a location where river water is moving past the side of the RST. The measurement should be recorded in milligrams per liter. Refer to dissolved oxygen meter manual for more information about the use, calibration and maintenance of the dissolved oxygen meter.

Water temperature (Celsius): water temperatures in the Stanislaus River will be measured with a YSI® 55 Dissolved Oxygen Meter. The location where water temperature is measured will be at a location where river water is moving past the side of the RST. The measurement should be recorded in Celsius units, and be recorded to the nearest tenth of a degree. Water temperature should be consistently taken around the same time every day.

Water turbidity (nephelometric turbidity units): turbidity levels in the Stanislaus River will be measured with an Oakton® T-100 Waterproof turbidity meter. Water samples should be collected from the river water moving past the side of the RST. When taking this sample, submerge the bottle entirely and allow it to fill completely (no air). Place the appropriate pre-labeled top for the trap location on the bottle from where the sample was taken. Place bottles into the cooler immediately and keep cool to preserve organics which may create added turbidity in sample. Take the bottles back to the office and quantify and record the turbidity on the Daily Trap Visit data sheet to the nearest tenth of a nephelometric turbidity unit (NTU). Refer to turbidity meter manual for more information about the use, calibration and maintenance of the turbidity meter.

Water velocity (meters per second): water velocities in the Stanislaus River will be measured with a Hach FH950 portable velocity meter. Measure the average water velocity in front of each screw trap, approximately halfway between the right pontoon and shaft and half the radius of the trap cone below the surface; record the value on the corresponding data sheet for that RST. Make sure the flow meter is programmed to present values in meters per second, be sure to re-zero the meter to measure the average velocity before taking the next reading, and record the average water velocity to the nearest tenth of a meter per second. Refer to water velocity meter for more information about the use, calibration and maintenance of the water velocity meter.

Weather: in two or three words, describe:

1. The amount of cloud cover (clear, partly cloudy, overcast, foggy)
2. If applicable, an indication of the amount of rain falling (sprinkle, rain, heavy rain).
3. If applicable, an indication of the amount of wind (slightly windy, windy)

APPENDIX D: Equipment Lists

Catch Visit Equipment List

The equipment that field biologists should take to the Stanislaus River trap site is as follows:

Clipboard	Trap Visit Data Sheets
Weight scale	Alka-Seltzer Gold
Knife	18-gallon tubs (2)
Surgical scissors	Pencils/Sharpies
Syringe	Fish ID book
Envelopes	Thermometer/dissolved oxygen meter
Stop watch (2)	Tools, screw drivers and crescent wrenches
First-aid kit	Nylon rope
Flashlights/headlamps	Zip ties
Rescue rope	Hub counter bolts/nuts
Pocketknife	Datasheets
WD-40	Paddles
Winch handle	Personal flotation devices
Waders	Water velocity meter
Wading boots	Digital camera
Ice chests	1/2 bucket for anesthesia bath
Measuring board	Scrub brushes (2)
Scoop nets (2)	Pitch fork
Dip net (1)	Aerator
Car battery	5 gallon buckets (8)
Cell phone	Vials for storing fish clips and whole fish
Salmon length-at-date chart	Vials for assessing water turbidity
200 proof ethanol	
Poly Aqua stress coat	

Bismark Brown Equipment List

Large tub	Marking Data Sheets
Large ice chest	Bismarck Brown Y stain powder
5-gallon buckets	Aerator
Frozen water bottles	Live cart

Anesthetizing Equipment List

Water	Container for mixing
Funnel	Latex gloves
1 liter container	Alka-Seltzer Gold

Elastomer Dye Equipment List

Clipboard
Syringe
Pencils
CO2 tank and regulator
3–5 Buckets
Aerator
Tie downs
Card table
Chairs
Net pen
Scoop net
Elastomer dye

Alka-Seltzer Gold
Marking Data Sheets
Thermometer
Deionized Water
Latex gloves
Live carts
Ice chests
Dip net
Towels
Poly Aqua Stress coat
Tool box

MadaJet toolbox with: Extra seals, Marine grease, Alcohol, Toothbrush, Dye powder, Inoculators, Dye and syringe.

APPENDIX E: Fish Species List for The Stanislaus River

Common Name	Species	Family	Status in Stanislaus River	Native/ Non-Native
Bigscale logperch	<i>Percina macrolepida</i>	Percidae	Possibly present	Non-native
Black bullhead	<i>Ameiurus melas</i>	Ictaluridae	Possibly present	Non-native
Black crappie	<i>Pomoxis nigromaculatus</i>	Centrarchidae	Possibly present	Non-native
Bluegill	<i>Lepomis macrochirus</i>	Centrarchidae	Present	Non-native
Brown bullhead	<i>Ameiurus nebulosus</i>	Ictaluridae	Possibly present	Non-native
Chinook salmon	<i>Oncorhynchus tshawytscha</i>	Salmonidae	Present	Native
Common carp	<i>Cyprinus carpio</i>	Cyprinidae	Present	Non-native
Golden shiner	<i>Notemigonus crysoleucas</i>	Cyprinidae	Present	Non-native
Goldfish	<i>Carassius auratus</i>	Cyprinidae	Present	Non-native
Green sunfish	<i>Lepomis cyanellus</i>	Centrarchidae	Present	Non-native
Hardhead	<i>Mylopharodon conocephalus</i>	Cyprinidae	Present	Native
Hitch	<i>Lavinia exilicauda</i>	Cyprinidae	Possibly present	Native
Inland silverside	<i>Menidia beryllina</i>	Atherinopsidae	Possibly present	Non-native
Largemouth bass	<i>Micropterus salmoides</i>	Centrarchidae	Present	Non-native
Pacific lamprey	<i>Entosphenus tridentatus</i>	Petromyzontidae	Present	Native
Prickly sculpin	<i>Cottus asper</i>	Cottidae	Possibly present	Native
Pumpkinseed	<i>Lepomis gibbosus</i>	Centrarchidae	Possibly present	Non-native
Redear sunfish	<i>Lepomis microlophus</i>	Centrarchidae	Present	Non-native
Riffle sculpin	<i>Cottus gulosus</i>	Cottidae	Possibly present	Native
River lamprey	<i>Lampetra ayresii</i>	Petromyzontidae	Possibly present	Native
Sacramento pikeminnow	<i>Ptychocheilus grandis</i>	Cyprinidae	Present	Native
Sacramento sucker	<i>Catostomus occidentalis</i>	Catostomidae	Present	Native
Smallmouth bass	<i>Micropterus dolomieu</i>	Centrarchidae	Present	Non-native
Splittail	<i>Pogonichthys macrolepidotus</i>	Cyprinidae	Possibly present	Native
Steelhead	<i>Oncorhynchus mykiss</i>	Salmonidae	Present	Native
Striped bass	<i>Morone saxatilis</i>	Moronidae	Possibly present	Non-native
Threadfin shad	<i>Dorosoma petenense</i>	Clupeidae	Present	Non-native
Threespine stickleback	<i>Gasterosteus aculeatus</i>	Gasterosteidae	Possibly present	Native
Tule perch	<i>Hysterocarpus traskii</i>	Embiotocidae	Present	Native
Wakasagi (Japanese smelt)	<i>Hypomesus nipponensis</i>	Osmeridae	Possibly present	Non-native
Warmouth	<i>Chaenobryttus gulosus</i>	Centrarchidae	Possibly present	Non-native
Western mosquitofish	<i>Gambusia affinis</i>	Poeciliidae	Present	Non-native
White catfish	<i>Ameiurus catus</i>	Ictaluridae	Possibly present	Non-native
White crappie	<i>Pomoxis annularis</i>	Centrarchidae	Possibly present	Non-native

Note: Other fish species may occur in the Stanislaus River. It should not be assumed the list above is a complete list.

The following references provide some of the information biologists can use to determine the species identification of unidentified fish in the Central Valley:

1. Moyle, P.B. 2002. Inland Fishes of California. University of California Press, Berkeley and Los Angeles, California, USA.
2. Wang, J.C.S. 2010. Fishes of the Sacramento-San Joaquin River Delta and Adjacent Waters, California: A Guide to Early Life Histories. Unpublished report prepared by the U.S. Department of the Interior, Bureau of Reclamation, Mid-Pacific Region and Denver Technical Service Center. Volume 44 – Special Publication. 411 pp.
http://www.usbr.gov/pmts/tech_services/tracy_research///tracyreports/TracyReportsVolume44r.pdf
3. Reyes, R.C., B.W. Bird, and P.F. Raquel. 2007. Guide to the Fishes of the Tracy Fish Collection Facility. Unpublished report prepared by the U.S. Department of the Interior, Bureau of Reclamation, Mid-Pacific Region and Denver Technical Service Center. Volume 36. 38 pp.
http://www.usbr.gov/pmts/tech_services/tracy_research///tracyreports/TracyReportsVolume36.pdf
4. [Http://www.usbr.gov/pmts/tech_services/tracy_research/photos/fish/ReyesFishGallery.html](http://www.usbr.gov/pmts/tech_services/tracy_research/photos/fish/ReyesFishGallery.html).
This website provides good photographs of some of the fishes in the Central Valley.

APPENDIX F: Key to Juvenile Chinook Salmon and Steelhead

From: Pacific States Marine Fisheries Commission
Smolt Monitoring Program
Guide To Fish Handling, Identification, And Condition
Revised April 2001

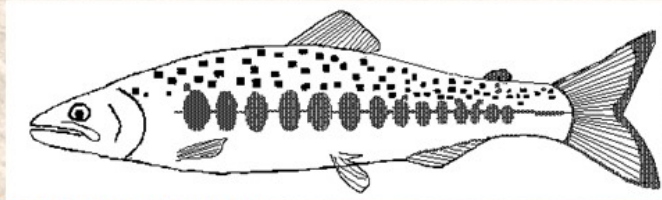


Figure 7. Distinguishing characteristics of Fall Chinook (*Oncorhynchus tshawytscha*).

1. Smaller eye that tends to turn down in head.
2. Deeper body, "football shape".
3. Usually more silvery in appearance.

Similar physical characteristics of Spring and Fall Chinook

1. Caudal fin forked, usually tipped in black.
2. Parr marks are large, vertically oblong, wider than the intervening spaces, and centered on the lateral line.
3. Anal fin rays are short, wedge shaped, and usually not pigmented.
4. Large, oblong spots on the back.

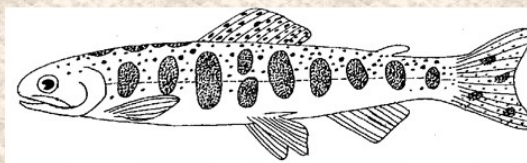
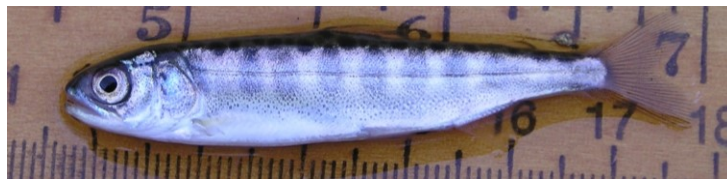


Figure 5. Distinguishing characteristics of Steelhead (*Oncorhynchus mykiss*).

1. Caudal fin not forked, with rounded lobes.
2. Parr marks nearly round, centered on lateral line.
3. Head more rounded than salmon when viewed from the top.
4. Dorsal fin has distinct black pigmented spots. In general, more spotting on fish.



APPENDIX G: Process for Staining Chinook Salmon With Bismark Brown Dye

Prepare Staining Solution at a Dosage of 1 packet of BBY (0.6 g) per ~20 gallons of water.

- a) Using a graduated cylinder, fill a large tub with 75 liters of water (approximately 20 gallons).
- b) Using a balance, measure out 0.6 gram of Bismarck Brown Y stain powder (this step should be prepared ahead of time in the lab).
- c) Thoroughly mix the Bismarck Brown Y powder in a small bottle with screw tight lid and shake to ensure powder is thoroughly mixed, and then add to the tub of water with fish to be stained.
- d) Place aerator and thermometer in tub.
- e) Keep water well oxygenated; use ice to maintain water temperatures within 2 °C of river water.

Immerse Salmon in the Stain Solution

- a) Count out fish to be stained with Bismarck Brown Y and place into dye solution.
- b) DO NOT anesthetize fish or add stress coat prior to immersion in dye solution.
- c) Record the number of fish placed in the stain solution.
- d) Set lid over tub to prevent fish from escaping and to protect fish from direct sunlight.
- e) Observe water temperature and fish activity regularly (every 5 to 10 minutes).
- f) Gently stir water while observing fish.

Monitor the fish, aeration, and water temperature in the tub during the staining period to detect signs of stress and possible causes. Fish will initially behave erratically; flare gills, and appear sluggish while in solution, this is normal. However, immediately remove individual fish displaying prolonged abnormal behavior and place into well-aerated recovery water.

Remove the Salmon From the Stain Solution

- a) Remove fish from the dye solution after two hours of immersion time in the solution and transfer them to a flow-through live cart to allow excess dye to drain out. Immediately submerge the live cart with lid in fresh river water to recover and wash out excess dye.
- b) Record the end time when fish were removed from the stain solution.
- c) Hold the stained salmon overnight. Prior to their release, remove and count all latent mortalities, and record the number of mortalities in the Mark and Release datasheet.

APPENDIX H: Process for Marking Chinook Salmon With Elastomer Dye

Fish can be marked with photonic dye using a needle-less photonic injector that places a small, semi-permanent colored elastomer dye mark between fin rays (Figure 1). Photonic dye marks are usually placed on the caudal fin for fry-size fish; however, the dorsal and anal fins can also be marked when fish are larger than 45 mm. Because photonic dye marks may last for several weeks (vs. the few days for fish stained with Bismark brown strain), and they provide the ability to provide unique batch marks that allow field staff to correlate the recapture of fish to a particular trap efficiency test (thereby drastically reducing the potential that that fish is assigned to the wrong trap efficiency release group), the juvenile salmon used that are marked for during trap efficiency test on the Stanislaus River will use either Bismark brown staining and photonic dye marks. This appendix describes the process for marking salmon with the photonic dye, and Appendix G provides the process for marking salmon with the Bismark brown strain. In some instances, fish may first be marked with photonic dye, and then stained with Bismarck brown stain.

Figure 1: Anal fin marked with needle-less gun using photonic pink dye.



A marking station is defined as the work space used by one biologist who is marking fish. Under normal circumstances, two biologists will mark fish, and a third biologist will record data and/or provide support to the two biologists marking fish.

Establish a work space

- a) Set up work station including table, chairs and canopy (if needed).
- b) Start a new Marking Data Sheet and record: Date, Project Location, Crew Member Names, Origin of Fish Stock (wild vs. hatchery), Release Code, Mark Applied, and record 100 fork lengths. Include life stage, and mortality for all 100 fork lengths taken.
- c) To prepare CO₂ tank, tap the bottom of tank against the ground softly 5 times and release a little CO₂ before attaching gun. This will clear out dry powder in CO₂ tanks which may damage the O-rings in marking gun. Then connect the regulator and marking gun.
- d) Attach marking dye hose and start CO₂ flow from the CO₂ tank.
- e) Place cutting board with marking tile into a shallow pan of water. Water level in pan should be about ¼ inch above the marking tile. Water level may be adjusted depending on each person's marking technique. Typically if water is too low, marking dye will

splatter when applied. If water is too high, marking may be inconsistent due to image refraction in the water.

- f) Fill a cooler 1/2 full with river or hatchery water, insert a working aerator in the tub, add Stress Coat to the cooler, then add up to 150 fish at a time to the cooler.
- g) Fill half a bucket with cool water and an appropriate dosage of Alka-Seltzer in the same way as stated earlier in this protocol.
- h) Fill recovery buckets about half full of river or hatchery water and add Stress Coat and place an aerator in the recovery bucket.
- i) Place about 10 to 20 fish per marking station in the Alka-Seltzer solution. Remember to keep in mind that you want to put as many fish in the solution as you can handle while you're marking fish. So for example, if you're a little bit slow going at marking at first you'll only want to place about 5-8 fish in the solution until you become faster at marking. The longer it takes you to mark a fish the longer the other fish sit in the anesthesia solution.

Mark Fish

- a) Take a random sample of 100 of the fish to be marked and measure the fork length and assess the life stage of each of those fish. Record those data on the backside of a Fish Marking Data Sheet.
- b) After all the fish have been measured and assessed, place fish on a plastic cutting board one at a time for marking after they've been anesthetized with the Alka-Seltzer solution.
- c) Photonic dye marks are usually placed on the caudal fin for fry-size fish; however, the dorsal and anal fins can also be marked when fish are larger than 45 millimeter in length.
- d) Apply the mark by starting with one pressure key turned out on the gun. Lightly place the gun tip onto the appropriate fin and pull the trigger. Be careful to not place the marking tip and mark too close to the body or fin margin as it can potentially injure fish.

Turn out one key at a time to increase gun pressure; test before marking.

If fin splits when marked, reduce the gun pressure or position.

Agitate the dye solution every couple of minutes, the microscopic elastomer beads will settle in solution quickly. If a marking becomes suddenly bolder, this is often times the solution. Dense concentration of elastomer beads can also clog marking gun and makes the remaining solution nearly useless when a majority of the beads are removed from solution.

- e) Count the marked fish, place them in recovery bucket, and record the “plus count” of marked fish on the data sheet.

Always check to ensure fish are recovering normally and have visible marks.

- f) If the gun jams, remove fish from the anesthesia solution before trying to fix jam. Guns can usually be fixed by running clean water through them or reversing the tip. NEVER put river water in the guns, they will clog! If this does not solve the problem after a few attempts, try using a different tip.
- g) When approximately 75 marked fish have accumulated in recovery bucket, transfer those fish to a live cart in the field or holding raceway in the hatchery where they will be held overnight.
- h) After 150 fish have been marked, develop a new mixture of Alka-Seltzer solution in 1/2 bucket to ensure that the solution maintains its potency, and continue to mark the fish until every individual has received a mark.
- i) After all fish have been marked, record the total number of fish marked on the datasheet. Mortalities should be recorded on datasheet and subtracted from total count.
- j) If fish are being held in river, carefully position live cart(s) in a secure location in river, secure the top and reinforce with a strap.

Clean up

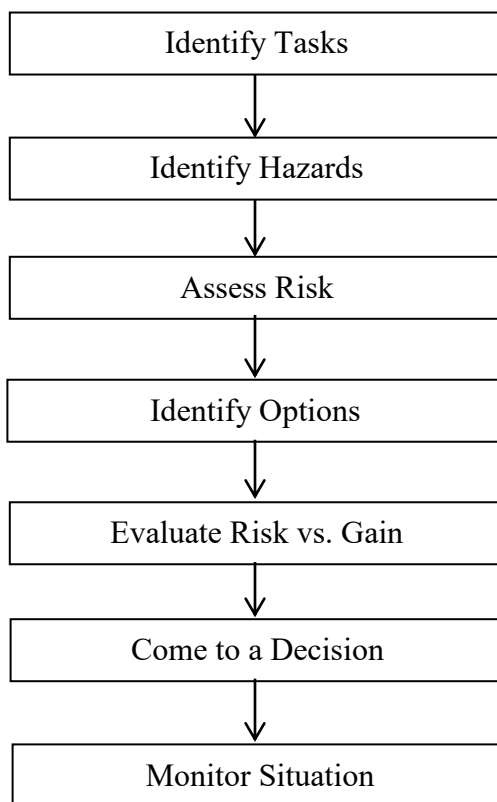
- a) Clean and load up all supplies. Marking guns should be cleaned thoroughly with deionized water and the cleaner solution provided to us by NewWest. Store the gun so it can dry in a short period of time. NEVER put a gun back into its case with dye in it.
- b) Field check data sheet(s) for completeness and accuracy.
- c) Make sure all the equipment is ready to be used again, and return all supplies to storage. Leave items out to dry thoroughly if they are still damp.
- d) Store the CO2 tank standing vertical.

APPENDIX I: Operational Procedure for Night Operations

Night time operations may be necessary in cases of increased river flow and/or major storm events where additional debris will likely be washed into the river and caught by the rotary screw traps. This debris can potentially cause the cones and live wells to be clogged with riparian/woody debris which can hinder trap operations. This debris can also lead to an increased mortality of salmonids being captured by the traps as they become trapped in the rotating cone instead of being directed to the live well. The traps can become clogged or blocked extremely quickly during these events. Because of this, a crew will go out during the evening and keep the traps clean and clear in order to reduce the potential for trap malfunction or fish mortality.

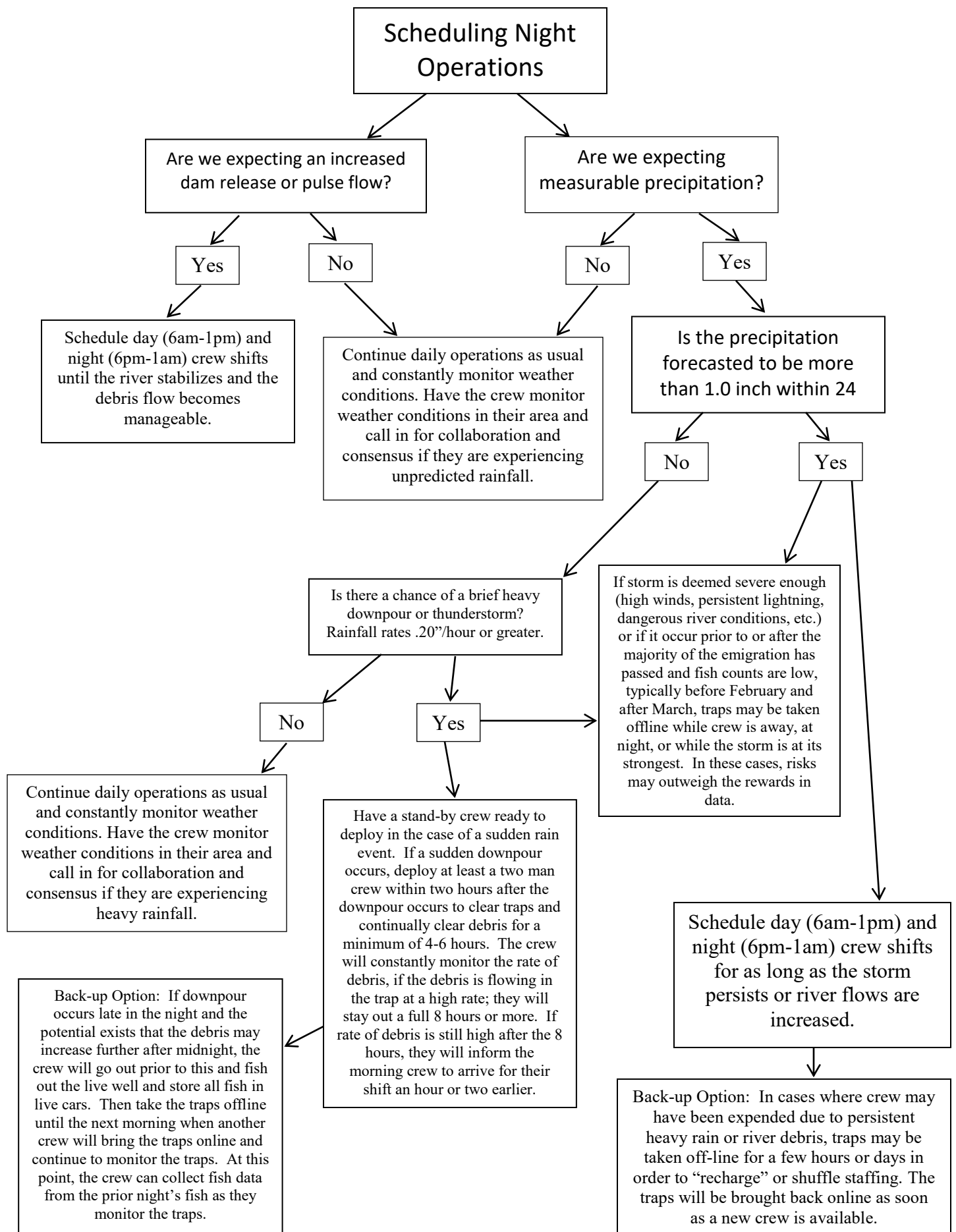
Night time trap operations will be considered on a case by case basis due to the inherent risk of operating in darkness and inclement weather. The decision will depend on the forecasted storm, amount of salmonids that may be passing the traps during this event and the potential hazards in regards to crew safety. A risk based assessment of these factors will be discussed with the field biologist, crew, and supervisors. Based on this discussion a determination will be made as to whether we believe a night time trap check is deemed warranted, unnecessary, or unsafe. The lead biologist will also determine the shift needed to perform the night time trap check depending on storm or forecasted river rise timing.

Steps for the decision making process regarding crew safety:



Hazards and risks that will be assessed:

- Storm severity
- Expected timing of storm
- Predicted river flow increase (from CDEC)
- Wind conditions
- Trap proximity to trees
- Amount of debris in river
- Types of debris in river
- Amount of fish expected to pass traps during event
- Experience of crew working on the trap
- Potential for injury
- Potential for equipment damage
- Likelihood of emergency crews reaching crew in case of emergency



Example Scenarios:

If storm precipitation is expected to increase river flow, but wind is expected to be less than 20mph, night operations will likely occur.

If storm precipitation and wind is likely to be high, but fish counts from the previous days have been less than 100 fish, night and trap operations may be suspended during the storm.

If storm is likely to have high winds (greater than 40mph), and traps are positioned underneath trees, safety of the crew may be compromised and trap operations may be suspended. If traps are positioned well away from trees, night operations will likely occur.

In instances where night operations will occur, a crew will go out to the traps in the evening and clear the live well and trap of debris. Fish will either be left in the live well or set aside in an additional live cart for assessment during the next day time trap check, depending on the quantity of fish. Fish assessment will be left for day time checks where there is enough natural light to accurately assess, measure, and weigh fish. If fish count for the night time check appear to be more than roughly 2,000-5,000 fish in the live well, those fish will be placed in a separate large live cart. A best judgment estimate will be used in these cases in order to prevent additional fish handling from counting individual fish. The ultimate goal of separating the fish is to prevent overcrowding inside the live well as more fish continue to be captured throughout the night. It is thought that an influx of fish will be flushed down the system during increased flow events. Additionally, most of the fish are captured late at night or in the early morning hours. These factors should be considered when deciding whether or not to leave fish in live well or store in live cart until the next daytime check.

During dam “pulse flows” only events, night time operations are likely to be performed by the crew. These are instances when dam operators will increase flows for a relatively short amount of time, typically less than a week, to help encourage the outward migration of salmonids from the Lower Stanislaus River into the San Joaquin River and delta. Typically the Bureau of Reclamation will give advanced notice of timing, duration, and magnitude of the pulse flow. Because of the importance of this data, special attention will be taken to ensure that the traps are operating during the entirety of the pulse flow. During these events, weather typically isn’t a factor.

Additional equipment for night operations:

- Navigation lights for jet-boat including the red/green side lights and white all-around light
- Deck lights for trap
- Deep Cycle battery to power deck lights
- Spot light
- Wearable head lights
- Flashlights
- PFD's with reflective tape and safety strobes
- US Coast Guard approved locator kit with flares
- Live carts for holding fish overnight