# Conservation Assessment of Steelhead Populations in Oregon 

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March 2001

## Table of Contents

Introduction ..... 1
Analytical Concepts and Methods ..... 2
Abundance Estimates ..... 2
Productivity ..... 3
Population Structure ..... 4
Conservation Thresholds ..... 5
Assessing Impact of Hatchery Fish ..... 6
Forecasting Persistence ..... 8
Assessment Results and Discussion ..... 19
Populations and Sub-populations ..... 19
Population Trends ..... 22
Observed Abundance and Conservation Thresholds ..... 22
Trends in Productivity ..... 24
Population Viability Analysis ..... 24
Assessment Synthesis ..... 26
Mortality Rate Assessment ..... 28
Hatchery Impacts ..... 32
Zones of Inference ..... 36
References ..... 37
Figures ..... 39
Appendices ..... 45
Appendix 1. Presumed steelhead populations ..... 45
Appendix 2. "Populations at a glance" ..... 47
Appendix 3. Summary of productivity estimates ..... 79

## Introduction

This report was prepared to accomplish two primary objectives. First, to provide updated information and analyses with respect to the conservation status of wild steelhead populations in Oregon. Secondly, it is intended to provide assistance to fish managers in their evaluation of the impact of steelhead fisheries and hatchery programs on the biological health of this species.

Both objectives have immediate application as ODFW is currently in the process of preparing a variety of management plans under the 4(d) provisions of the Endangered Species Act (ESA) at the request of the National Marine Fisheries Service (NMFS). Although this effort is focused on listed populations in the Columbia Basin, similar ESA issues potentially exist for coastal steelhead populations. For example, as a result of a recent court order, NMFS is once again evaluating steelhead populations of southern Oregon and northern California for possible addition to the threatened species list.

The report is organized by data sets that are thought, in most cases, to represent demographically independent groups of naturally reproducing steelhead following the concepts described by McElhaney et al. (2000). However, one reality of these data sets is that they are not evenly distributed throughout Oregon. Depending on the presumed population structure and interrelationships of these populations, there are invariably natural production units for which data does not exist. As a result, to assess this species it is necessary to utilize a "zone of inference" around each data set. In some cases this zone encompasses only a single population. However, in other situations, it includes multiple populations.

With respect to a status assessment, the construction of these zones of inference has more practical significance than does population and subpopulation structure. For example, if the picture from multiple monitoring sites within a given region appears uniformly healthy, then it seems reasonable to conclude that the steelhead within this region are healthy, regardless of which population or sub-population they belong to. However, if the picture is not uniform, then a single regional zone of inference becomes difficult to justify. Indeed, in this latter case the status of steelhead in unmonitored production areas may be impossible to resolve with any certainty. Especially if there is no knowledge as to which population they might demographically cluster with. The
conservative approach to such a situation is to assume the status of these unknown populations is bad.

Therefore, the resolution of these zones of inference is the crux in assessing Oregon's steelhead populations. The methodologies used in this report were tailored to address this analytical problem. As a result, they produced an assessment with more emphasis on among population comparisons as opposed to descriptive treatments of individual populations.

Finally, a word about the organization of this report. The results and discussion section covers population and sub-population descriptions, recent trends and abundance, an estimate of critical conservation thresholds for each population with respect to abundance, trends in productivity, a population viability analyses for each population, and a synthesis of these results. In addition, three supplemental sections area also included. The first describes an evaluation of how sensitive the conservation status of each population is to changes that cause an increase in mortality (for example fishing mortality). The second supplemental section examines the potential impact of naturally spawning hatchery fish on the productivity of natural populations. The last section is a brief comparative summary of report findings with particular attention to the zones of inference for which they apply.

## Analytical Concepts and Methods

## Abundance Estimates

For each monitoring location, annual estimates of adult spawner abundance or density (fish per mile) were determined from direct adult enumeration at counting facilities (Rogue, Umpqua, Clackamas, Sandy, Hood, and Umatilla populations), from redd counts (most other locations), or from mark-recapture population estimate techniques (Deschutes and Walla Walla populations). Conversion of redds per mile to spawners per mile, discrimination between hatchery and wild fish, and estimation of cumulative fishery mortality on wild steelhead was similar to methods described by Chilcote (1998). Estimates of pre-harvest abundance for wild steelhead were obtained by dividing annual estimates of spawner abundance by 1 minus the associated harvest rate.

## Productivity

Productivity, as used in this report, is the number of adult offspring (recruits) produced per spawner. It is determined by counting all of the fish that spawn in a monitoring area (both hatchery and wild fish) and dividing this number into the number of pre-harvest offspring produced by these spawners. Steelhead have a complex life history with multiple ages of return and the capacity to spawn more than once. Therefore, estimating recruits is process of apportioning each year's return into the correct parental brood year and then obtaining a brood year total my adding up its apportioned amount across multiple return years.

Productivity, in its various expressions, is probably the most important factor to consider in assessing the conservation status of a species. It is related to the innate ability of a population to rebuild its self and therefore relates directly to forecasting the persistence of the population. In addition, the incorporation of underlying trends or cycles in productivity is often critical in understanding the true biological health of a population. Further, by regressing observed recruits per spawner and total spawner abundance it is possible to estimate the capacity of a given habitat to produce adult steelhead. In other words, it is possible to estimate the number of spawners needed to seed the available habitat to maximum production (maximum seeding).

For most naturally reproducing populations, productivity (recruits per spawner) decreases as the spawner abundance increases. This is because as juveniles fill up the available habitat, the proportion that is able to survive becomes less and less. Therefore, to estimate the productivity of a population in a consistent manner, it is necessary to standardize the recruit per spawner data with respect to spawner abundance. In this report, this standardization process was accomplished by estimating the a parameter of the Ricker recruitment equation,

$$
\text { Recruits }=\text { Spawners(2.718(a+B(Spawners)) }
$$

Equation 1
In this recruitment relationship both the $\boldsymbol{a}$ and $\mathbb{B}$ parameters were estimated using the linear regression method, where the general equation, $\mathrm{y}=\boldsymbol{a}+\mathbb{B}(\mathrm{x})$, was transformed to:

$$
\operatorname{Ln}(\text { Recruits } / \text { Spawner })=\boldsymbol{a}+\mathbb{B}(\text { Spawners })
$$

Equation 2
Therefore, for a data set of paired observations of spawner abundance and $\operatorname{Ln}$ (recruits/spawner), the $\boldsymbol{a}$ parameter is the $y$-intercept and the $\mathbb{B}$ parameter the regression line slope, which is almost always negative.

Because these values remain the same for a population, regardless of its spawner abundance, they serve as a standardized way to compare different populations and as a way to compare the same population at different intervals of time. The a parameter serves as a means to compare population productivity. In addition, the inverse of the $\boldsymbol{B}$ parameter, $1 / \boldsymbol{B}$, can be shown to be the spawner abundance which generates the maximum number of recruits. Therefore, the $\boldsymbol{B}$ parameter can be used to estimate how many recruits a population can produce, while the a parameter estimates how efficient the population is in producing them. For the purposes of this report, $1 / \boldsymbol{B}$, the number of spawners necessary to achieve maximum production of recruits, will be referred to as "maximum seeding".

Estimates for $\boldsymbol{a}$ and $\mathbb{B}$ were generated for 27 of the 31 data sets examined. The four data sets omitted were from relatively new monitoring sites and as such did not yet have a sufficient number of data points to estimate their recruitment parameters. Most data sets examined extended back to 1974. Rather than fitting a single recruitment curve to all of the data from each site, a series of multiple curves, and associated estimates of $\mathfrak{a}$ and $\mathbb{B}$ were determined for each data set. This was done to examine the temporal variation in productivity.

These multiple curves were build upon a moving 7-year sequence of spawner/recruit data. For example, for a population having spawner/recruit data beginning in 1974, the first recruitment curve was estimated for the spawners of 1974 to 1980 and their subsequent recruits. The next recruitment curve was based upon the production of 1975 to 1981 spawners. The third curve, for 1976 to 1982 spawners and so forth until the end of the data set. Depending on the length of the data set, 10 to 25 recruitment curves and associated values for the Ricker equation parameters $\mathbb{a}$ and $\mathbb{B}$ were generated for each population.

## Population Structure

As stated earlier, the majority of the data sets presented in this report were thought to represent demographically independent populations of naturally reproducing steelhead. In general the boundaries for these populations followed those described by Kostow et al. (1995). However, in the interest of meeting the test of demographic independence, steelhead from several smaller populations were lumped together as one. This was done largely on the basis of geographic proximity and relative size of each watershed. Although somewhat logical, the empirical evidence to justify such "lumping" was generally lacking. This is because data is rarely collected from steelhead returning to these smaller basins.

Sub-populations were also identified in this exercise, largely at the suggestion of biologists familiar with the local area. The criteria used for identification of these sub-populations were known discontinuities in hydrology, elevation, geology, temperature regime, vegetative cover, basin aspect, and spawn timing. It was assumed that these physical differences were capable of causing some degree of reproductive isolation and genetic adaptation. In addition, a limited set of biochemical data was used to gauge the level at which divisions likely occurred.

In the case of many populations, especially in the Columbia basin, sufficient data were available to gauge their likelihood of demographic independence. This took the form of comparing trends in abundance, productivity and relative vulnerabilities to extinction.

## Conservation Thresholds

To provide better a context for estimates of spawner abundance, two numerical conservation thresholds were developed, "critical" and "viable". These thresholds were intended to be ODFW's interpretation of the critical and viable thresholds described by McElhaney et al. (2000) and NMFS (2000). As used in this report, these thresholds represent one of several biological criteria used to determine the overall status of steelhead populations in Oregon. In addition to these conservation thresholds, natural production benchmarks representing 50\% and $100 \%$ of maximum seeding were developed.

Several of the populations examined contain naturally spawning hatchery fish. However in determining the conservation thresholds for these populations, the reproductive contribution of these hatchery fish was excluded. This approach was taken to ensure that the conservation thresholds would represent the natural, self-sustaining response of populations to critical levels of abundance in consistent manner. Hatchery fish, when they are present, are sources of reproductive effort whose origins are essentially external to the natural population. Potentially, this can give the illusion that a population is self-sustaining at low levels, when in fact this is not the case. Therefore, hatchery fish were excluded from the conservation threshold estimation procedure in order to achieve a standardized means of describing these thresholds.

The approach for estimating population specific values for the viable threshold and the other 2 natural production benchmarks was largely the same. As discussed previously, multiple recruitment curves were calculated for each population. Each of these curves had an associated value for the $\boldsymbol{B}$ parameter. For each population, these $\boldsymbol{B}$ values were
averaged and then divided into 1 to obtain an estimate of the average number of spawners needed to produce maximum recruitment. The natural production benchmarks were calculated as $0.5 / \boldsymbol{B}$ and $1.0 / \boldsymbol{B}$, corresponding with $50 \%$ and $100 \%$ of the number of spawners necessary for maximum recruitment (maximum seeding).

The viable threshold was set at $20 \%$ of the maximum seeding level or $0.20 / \boldsymbol{B}$. The logic for selecting $20 \%$ of $1 / \boldsymbol{B}$ as the threshold was based upon the lack of confidence in predicting the response of populations at escapement levels less than this level. The primary reason for this uncertainty was that escapements below these levels have rarely been observed in the data sets. Averaged across all populations and years, only $6 \%$ of the spawner escapement data points were less than $0.20 / \boldsymbol{B}$. Therefore, very little information was available to investigate how these populations actually performed at low escapement levels. In light of these shortcomings, it seemed logical that this threshold of uncertainty would suffice as the viable threshold.

To examine the consistency with concepts described by McElaney et al. (2000), viable thresholds for each population were evaluated to determine if they represented the boundary above which the probability of extinction in 100 years was "negligible". Using criteria presented by Allendorf et al. (1997), extinction risk was considered "negligible" if the probability of population extinction in 100 years was $5 \%$ or less. A population viability assessment (PVA) model, described in subsequent sections of this report, was developed and used to make this evaluation.

The critical abundance level for each population was determined directly from the PVA model. In the context of PVA models, Mace and Lande (1991) proposed the following standard for endangerment: a $20 \%$ probability of extinction over a period of 10 generations. For the purposes of this report, their classification of "endangerment" was assumed to be synonymous with "critical". Adopting this standard, the critical abundance threshold was defined as the number of spawners, that if left alone to naturally reproduce for 50 years (approximately 10 generations) would result in the extinction of the population more than $20 \%$ of the time. This critical abundance was estimated for each population by seeding each PVA model run with fewer and fewer initial spawners until a $20 \%$ extinction probability was achieved.

## Assessing the Impact of Hatchery Fish on Natural Production

To varying degrees, hatchery fish were present in nearly half of the populations examined. There are numerous studies suggesting that naturally spawning hatchery fish may be less successful at leaving
surviving offspring than are wild fish (Nickelson et al. 1986, Reisenbichler and McIntrye 1977, Chilcote et al. 1986, Leider et al, 1990, and Reisenbichler and Rubin, 1999). However, whether or not these observations apply universally to all naturally spawning hatchery fish under all conditions remains unknown. This question had immediate implications to the assessment of steelhead populations in Oregon. Specifically, for the populations examined, did the productivity of naturally spawning hatchery fish and wild fish differ and by how much? If differences exist, then this element would have to be incorporated into the assessment. Complicating matters further, the past and future presence of hatchery fish in Oregon's steelhead populations is not static. Changes have occurred in the recent past and are expected to occur in the near future.

In light of these considerations, it was necessary to determine the likely impact of hatchery fish on the productivity of these naturally reproducing populations. If following this determination it was found that the impact of hatchery fish could be significant, then there would be a need to add this factor to the assessment protocol. Further, in considering this factor, it would be necessary to incorporate the known dynamic nature of past, present, and future hatchery programs.

To assess the potential impact of hatchery spawners on natural production, overall population productivity was compared to the relative abundance of hatchery and wild fish on the spawning grounds. It was hypothesized that if hatchery fish had an adverse impact on natural production, then populations with high proportions of hatchery fish would be less productive than populations where the proportion of hatchery fish was low.

Values for the Ricker a parameter were used as an index of productivity. As described earlier, multiple recruitment curves, based upon 7 -year spawner sequences, were fit to each population's data. This generated a time series of a parameter values for each population. The average proportion of hatchery fish for each 7 -year sequence was matched up with the corresponding value estimated for the a parameter. Data from all populations was prepared in this fashion and grouped according to time period, beginning with the 7 -year sequence from 1978 to 1984, and ending with the 1989-95 sequence. This resulted in population productivity and hatchery proportion data groups for 12 time intervals.

For a majority of these populations the proportion of hatchery fish was 0.00 . However, three of these populations without hatchery fish were included in this evaluation: Joseph, Lower NF John Day, and North Umpqua winter steelhead. The remaining populations used in this analysis all had a history of naturally spawning hatchery spawners.

These included the following populations: Imnaha, Upper Grande Ronde, Umatilla, Deschutes, Sandy, Clackamas, Sandy, Molalla, North Santiam, South Santiam, North Umpqua summer steelhead, Rogue summer steelhead, and Rogue winter steelhead.

For each 7-year time period, a linear regression was performed, with ' $x$ ' being the proportion of hatchery fish and ' $y$ ' the a parameter value. Represented by the equation:

$$
\mathrm{y}=\text { Slope }(\mathrm{x})+\text { Intercept }
$$

Equation 3
If, in reviewing the results of these analyses for the 12 time intervals, regressions were found to be statistically significant and having negative slopes, then it was to be concluded that hatchery fish were less productive than wild fish.

## Forecasting Persistence

A key component of this assessment was forecasting the likelihood of extinction for each population under a variety of possible future conditions. The status of populations whose probability of extinction was forecast to be low or non-existent was assumed to be relatively healthy. In contrast, when the probability of extinction was high, the population was assumed to be at risk and unhealthy. To make these forecasts, a population viability assessment (PVA) model was developed. PVA models are frequently used in conservation biology to assess the vulnerability of populations to extinction. Such models have several common elements.

First, to forecast the abundance of a naturally reproducing population at some point in the future (e.g. 100 years) it is necessary to select an expected reproductive rate for this time period. Usually, it is assumed that the reproductive rate in this future period will be similar to the rate observed in the recent past. In the model used here, the reproductive rate that is assumed in the future was not a single number derived from averaging past rates observed for each population in recent years.

To provide a more realistic model of long-term population recruitment, a different rate is selected for each generation of recruits forecast. These reproductive rates are randomly selected from a pool of possible values. This pool of possible values is usually generated on the basis of the variation in reproductive rates observed for each population.

As an outcome of randomly selecting reproductive rates, the population abundance at the end of each cycle of a model run will not be the same.

Therefore, another characteristic of PVAs is that multiple forecasts (usually between 500 and 5000) are made for each set of conditions tested. The number of these forecasts where the population is less than a numerical threshold are then counted and divided by the total number of forecasts made. The result is an estimate of the probability that a population will be less than a numerical threshold within the test period. Often the numerical threshold is set at zero, or extinction. Therefore, the results of a PVA are usually stated in terms of the probability of extinction at some future point in time.

Finally, the interpretation of these PVA results requires a consistent standard for how much risk is acceptable. For example, a commonly used standard is that a population is endangered if the probability of extinction is greater than $20 \%$ at a point in time 10 generations into the future (Mace and Lande, 1991). Likewise, if a population has a probability of extinction of 5\% or greater after 100 years, then it commonly would qualify for a threatened classification (Allendorf et al. 1997 and Thompson, 1991).

The specifics for the PVA used to evaluate the steelhead populations considered in this review were as follows. The natural spawning abundance (or density) of hatchery plus wild fish for the most recent 6 years (1995 to 2000) were used to seed each PVA model run. Once seeded, the recruits from this escapement were forecast, subjected to a hypothetical fishery and then converted to spawners from whom the next generation of recruits was forecast. This process was repeated until a period of either 50 or 104 years had passed. Upon completion, the number of spawners in the last 6 years of the forecast period was examined to determine if they had declined to the zero. If zeros were forecast for all six ending years, then the cycle was recorded as an extinction event. Once completed, a second cycle was started using the same number of initial spawners and model conditions. The same forecasting process was run forward for the same length of time. If a second extinction event occurred it also was recorded. A single model run consisted of 1000 of these repeated cycles. The probability of extinction was calculated by dividing the number of cycles that resulted in an extinction event, by 1,000 , the total number of cycles for each model run.

In addition to model runs to estimate the probability of extinction, the PVA model was also used to estimate the probability that a population would decline to levels below its viability abundance threshold. In this case, if the forecast abundance of the population were less than the viable threshold in the last 6 years of the model run, then this would be recorded as a "viability risk event".

To estimate the number of recruits within each iteration of the model run it was necessary to develop a method to randomly sample reproductive rates from a pool of values that could be reasonably expected to occur in the future. It was assumed that the magnitude and pattern of reproductive rates observed for these populations over the last 25 to 30 years could be used to approximate future reproductive rates. Obviously, this assumption could not be tested directly. Of special concern is the possibility that future reproductive rates will be less than those of the past. If this were to occur, the PVA results would overestimate the likely persistence of these populations. Therefore, to reduce the risk of over-estimating persistence, the average survival of rates used in the model runs were $30 \%$ from what had actually been observed in the recent past.

The multiple recruitment curves fit for each population formed the basis from which the future reproductive rates were determined. As discussed earlier these curves were based upon a moving, 7-year sequence of spawner and recruit data. Each time period was assigned to a code year. The code year was the mid-point of the brood year sequence from which each recruitment curve was generated. For example, the code year 1972 represented the recruitment function for the fish that spawned from 1969 to 1975 . Code year 1973 indexed the recruitment function for fish from 1970 to 1976. Associated with each code year was a value for the a and $\mathbb{B}$ recruitment curve parameters.

A preliminary examination of the a parameter values for the code years from 1972 to 1993, suggested that cyclic pattern existed for most steelhead populations. It appeared that a peak in a values occurred with the code year 1981, followed by a low point in a values with the 1990 code year, a time frame of 9 years. Further, that 9 years prior to the peak in 1981, there appeared an earlier low point corresponding with the 1972 code year. This pattern suggested a symmetrical fluctuation of a parameter values, having a period of approximately 18 years. It was hypothesized that this cyclic pattern was a reasonable model for what would occur in future years. Therefore, this cycle was extended forward to the code year 2120. For each population, observed $\boldsymbol{a}$ and $\boldsymbol{B}$ values for code years 1972 to 1992 were repeated in reverse and then forward sequences such that they tracked the assumed cycle from code year 1994 to 2120. These "dummy data" were the basis from which the reproductive rates used in the PVA model were drawn.

As each cycle of the model run proceeded, the number of recruits would be calculated for each code year and then these recruits assigned to different future years of adult return depending on the average age distribution of the population. For example, if the recruitment from spawners in 2030 was 1,000 fish and the average age distribution was
$10 \%$ age $3,60 \%$ age 4 , and $30 \%$ age 5 , then 100 fish would be assigned to the pool of potential spawners of 2033, 600 to the year 2034, and 300 fish to the year 2035. The spawner code year would then be advanced one to the year 2031 and a new set of recruits produced and distributed to the appropriate future brood years. In this way, the number of spawners was build up for each future generation in the model. Obviously, when a fishery mortality rate was imposed, there was an intermediate step of removing some the recruits before they could be classified as spawners.

However, estimating the recruits from any one code year of spawners was not a simple deterministic process of using the Ricker recruitment formula with the specific values for $\boldsymbol{a}$ and $\boldsymbol{B}$ assigned to each code year. To introduce randomness into the process, the $\boldsymbol{a}$ and $\boldsymbol{B}$ values were drawn from a 7 -year sequence of code years that included the reference code year as its midpoint. For example, for estimating recruitment for the fish that spawned in 2023, a code year from 2020 to 2026 was randomly selected. Once selected, the associated values for the $\boldsymbol{a}$ and $\mathbb{B}$ parameters were used in the Ricker function from which the recruits for the 2023 brood year were calculated. The decision to use a sequence of 7 years to randomly draw the recruitment parameters for the each calculation was a compromise. In picking a time interval for this purpose, it was necessary to have enough data points to ensure that a reasonable degree of randomness would be introduced into the PVA model. However, this had to be balanced with not having so many data points that the cyclic nature of the underlying reproduction rate would be homogenized. For example if the recruitment parameters were drawn randomly from an 18 -year sequence of code years, the cyclic pattern (which appears to have a period of 18 years) would be effectively lost from the PVA simulation.

There were four additional, yet significant, nuances placed upon the process of forecasting recruits within the PVA model. First, as noted earlier, it was unknown if the relative survival rates observed over the last 20 to 30 years would be the similar to those of the future. It is possible that these future rates will be lower. Persistence forecasts based on higher than realized reproductive rates will yield extinction risks that are too low. Therefore, in order to make this outcome less likely, the survival rates for all model runs were assumed to be $30 \%$ less than those actually observed in the last 30 years. To make this adjustment, the number of recruits calculated each time the Ricker recruitment function was used within the model was reduced by $30 \%$.

The second nuance dealt with forecasting recruits when the number of spawners was very large. The nature of the Ricker recruitment function is such that once spawner escapements exceed $1 / \mathbb{B}$, the level necessary
for maximum recruitment, the number recruits begins to decline. In some cases this rate of decrease may be quite rapid. Although, the number of populations for which spawner escapements in this upper range have been observed is limited, a preliminary assessment of their subsequent recruitment did not appear consistent with a recruitment function that had a strongly declining right-hand limb. Indeed, alternate recruitment functions that do not have this decreasing recruitment behavior at high escapement levels, such as the Beverton-Holt equation, are often used for species like steelhead (Burgman, et al. 1993). However, the Beverton-Holt model was not used in the present population assessment for a variety of reasons. The primary reason being that the purpose of this assessment was to determine the risk of extinction for these populations. Therefore, the portion of the recruitment function that was of greatest importance was the performance at spawner densities considerably less than $1 / \boldsymbol{B}$. In this range the Ricker function appears to provide as good, if not better, fit to the pattern of recruitment in steelhead than the Beverton-Holt model.

However, to use the Ricker equation for the PVA model, a modification was necessary in order to get around the inherent problems when spawner density was high. To prevent the tendency for the Ricker function to underestimate recruits at these high spawner densities, a conditional step was added to the recruitment forecast process. This conditional step was triggered when the simulation model produced a number of spawners greater than $1 / B$. Under this circumstance the program reset the number of spawners to $1 / \boldsymbol{B}$. This step essentially meant that for any spawner escapements greater than necessary for maximum production, the recruits would be equal to maximum production. This eliminated the descending right-hand limb of the Ricker recruitment function, which for steelhead was felt to be problematical.

Another feature added to the recruit forecasting process was a modification to the model when spawner densities were very low. The available data sets contained only a few points from extremely low escapement levels. Therefore, it was not clear how the recruitment process actually functioned at these low levels. As noted earlier, this sense of uncertainty lead to the designation of a viable population threshold for spawner levels less than $20 \%$ of $1 / \mathbb{B}$. For spawner escapements less than $0.20 / \mathrm{B}$, it was difficult to confirm that the Ricker function was a good representation of the recruitment process. Of particular concern was the chance that at these low levels the expected recruitment mechanisms may begin to fail (Glipin and Soule, 1986). Either because of genetic problems or the inability of spawners to find mates in a low-density environment, the productive capacity of a population may decrease as the population declines below some critical
level of spawners. If such factors come into play for depressed steelhead populations, then using an unmodified Ricker model to forecast recruitment would overestimate a population's resiliency and productivity. This could lead to overly optimistic conclusions about the resistance of the population to extinction. Since there was a strong desire to avoid this type of error, the recruitment function was modified to be less productive at low spawner densities. Although there was no empirical evidence from which to base this modification, it was added because from a conservation management standpoint, the consequences of overestimating the probability of extinction were more acceptable than those associated with underestimating the probability of extinction.

The specific depensation modification to the recruitment function at low spawner densities was as follows. First, it was assumed that for spawner densities less than $0.04 / \mathbb{B}$, the population was essentially extinct and so the number of recruits produced from escapements less than $0.04 / \boldsymbol{B}$ was set to equal zero. For spawner densities in the range from $0.04 / \boldsymbol{B}$ to $0.20 / \boldsymbol{B}$, the recruitment function was a simple linear relationship beginning with zero recruits when spawner density was $0.04 / \mathbb{B}$, and increasing proportionately to the number of recruits forecast by the Ricker function when the spawner levels was $0.20 / \mathbb{B}$. For example, suppose for a particular population the levels of $0.04 / \mathrm{B}$ and $0.20 / \mathrm{B}$ were estimated to be 60 and 300 spawners, respectively. Using this approach, the number recruits from an escapement of 60 fish would be set at zero. The number of recruits from an escapement of 300 spawners, as predicted by the Ricker recruitment function would be suppose 600 fish. Therefore, the recruitment from an intermediate escapement, suppose 90 fish, would be calculated as: $600[(90-60 /(300-60)]=600(0.125)=75$ recruits.

Lastly, and the most complicated addition to the recruitment forecasting process was the mechanism devised to account for differences in reproductive success between naturally spawning hatchery and wild fish. As discussed earlier, studies have shown that at least in some situations, the reproductive success of naturally spawning hatchery fish is much less than it is for wild fish. Further, methods were also described earlier to determine if evidence for this difference existed for the steelhead populations examined by this assessment. Because, it was thought likely that such differences would be found, an approach for making adjustments to the recruitment forecasting process of the PVA model was developed.

Obviously, for populations that have never been exposed to hatchery fish and were assumed to remain in this condition for the future, no adjustments were necessary. In particular, the reproductive rates for the base period (1972 to 1992 code years) could be projected forward into
future without modification. Likewise, adjustments were not necessary for populations in which the proportion of hatchery fish had remained relatively stable through the base period and was expected to remain unchanged in the future.

However, the populations that were a potential problem were those for which had considerable fluctuation in the proportion of hatchery spawners during the base period and for which it was also unlikely this pattern would be repeated in the future. Also a problem, were those populations for which the average proportion of hatchery spawners in the future was expected to differ from the proportion during the base line time period. For example, if the proportion of hatchery fish in the base period was in the range of 30 to $50 \%$, but was expected to decline to $10 \%$ in the future, then using the reproductive rates observed during the base period to forecast the recruitment in future years would underestimate the productivity of the population and its resistance to extinction. (Obviously, this problem would only exist if a difference in productivity between hatchery and wild fish belonging to the populations evaluated in this assessment were confirmed). Likewise, the future productivity and resistance to extinction would be overestimated for a population that had very few hatchery fish during the base period, but was expected to have a much higher proportion in the future. The approach for correcting these potential sources of error was as follows. First, the theoretical relationship between the overall productivity of a population and the proportion of hatchery fish in the population was represented by:

$$
\boldsymbol{a}=\mathrm{P}_{\mathrm{w}}\left(\boldsymbol{a}_{\text {wild }}\right)+\mathrm{P}_{\mathrm{h}}\left(\boldsymbol{a}_{\text {hatchery }}\right) \quad \text { Equation } 4
$$

where $a$ is the Ricker recruitment parameter calculated for the population at a particular time interval, $\mathrm{P}_{\mathrm{w}}$ and $\mathrm{P}_{\mathrm{h}}$ are the respective proportions of wild and hatchery fish in the natural spawning population, $a_{\text {wild }}$ is the recruitment parameter that would have been estimated for this population were the only spawners wild fish, and $a_{\text {hatchery }}$ the recruitment parameter for a spawning population consisting only of hatchery fish.

If the values for $a_{\text {wild }}$ and $a_{\text {hatchery }}$ and the future proportion of hatchery spawners could be known then the overall productivity of the population, a, could be calculated. The past and present proportion of hatchery fish can be resolved with relative ease. However, the sequence of a values for the base period (1972 to 1992 code years), are overall measurements and do not contain separate estimates for $a_{\text {wild }}$ and anhery. Therefore, a method was needed to estimate awild and anatchery for the base period so that these values could be used to compute a more realistic overall population value for a in future years under scenarios where the
proportion of hatchery fish was expected to change. Equation 4, discussed previously, provides a means to do this.

To evaluate the potential for a relationship between the proportion of hatchery fish and overall population productivity, a linear equation was fit to the paired data sets from 15 steelhead populations for each of 12 time periods evaluated (code years 1981 to 1992) (see the methods section "Assessing the Impact of Hatchery Fish on Natural Production"). Slope and intercept parameters for a linear regression was estimated for each of the 12 time periods. Based upon each regression equation, theoretical $\boldsymbol{a}$ values for a population comprised entirely of wild fish and a second hypothetical population comprised entirely of hatchery fish were calculated (i.e., x in the regression equation was set to 1.0 to obtain the upper range limit for $\boldsymbol{a}$ and set to 0.0 to obtain lower range limit for $\mathfrak{a}$ ). This calculation was made for all 12 regressions and resulted in theoretical $\boldsymbol{a}$ values for a population comprised of $100 \%$ wild fish and one comprised of $100 \%$ hatchery fish for each of the time intervals corresponding to code years 1981 through 1992.

For the purposes of the PVA, this generalized model for assessing differences between hatchery and wild fish in terms of productivity had to be further modified so that it was specific to each population. This was accomplished in the following manner. First, the average proportion of hatchery fish for each population for each code year from 1981 to 1992 was calculated. Recall, that the estimates of $\boldsymbol{a}$ and $\boldsymbol{B}$ for each code year correspond with the recruitment from 7 brood years of spawners. Therefore, 7 years of data were used to compute each average.

Average hatchery fish proportions were then substituted into the generalized hatchery-wild regression models corresponding with the same time interval and an expected overall $\boldsymbol{a}$ value was calculated. For example, if the observed average proportion of hatchery fish for code year 1985 was 0.30 , then this value would be substituted for x in the generalized regression equation corresponding to the code year 1985 (i.e., 1982 to 1988 brood years). This step yielded an expected overall population value for productivity, $a_{\text {exp. }}$. This expected value was compared to the a value actually calculated for the specific population and used to standardize the estimates of productivity for wild fish and hatchery fish as shown in Equations 5 and 6.

$$
\begin{array}{cc}
\boldsymbol{a}_{\text {pop_wild }}=\boldsymbol{a}_{\text {wild }}+\left(\boldsymbol{a}_{\text {obs }}-\boldsymbol{a}_{\text {exp }}\right) & \text { Equation } 5 \\
\text { and } \boldsymbol{a}_{\text {pop_hatchery }}=\boldsymbol{a}_{\text {hatchery }}+\left(\boldsymbol{a}_{\text {obs }}-\boldsymbol{a}_{\text {exp }}\right) & \text { Equation } 6
\end{array}
$$

Where $a_{\text {pop_wild }}$ is a standardized estimate of productivity for the wild fish that spawned in this specific population during a specific time interval;
$\boldsymbol{a}_{\text {wild }}$ is the estimated productivity for the same time interval based upon the generalized model for differences between hatchery and wild fish developed from regression analysis of 15 populations as described earlier; $\boldsymbol{a}_{0 b s}$ is overall population productivity actually observed for the population during this specific time interval: and $a_{\text {exp }}$ is the expected overall population productivity for this time interval as estimated from the generalized hatchery-wild regression model. The terms in Equation 6 are similar to Equation 5, except that in the case of the former they refer to hatchery fish.

Once population specific estimates for $\boldsymbol{a}_{\text {pop_wild }}$ and $\boldsymbol{a}_{\text {pop_hatchery }}$ were obtained for all years in the base period, they were expanded to the future code years (1995 to 2120). This expansion was done by repeating the sequence of values for the base period in reverse and forward order as necessary to follow the presumed 18-year productivity cycle.

Once this "dummy data" had been entered, the simulated recruitment for each code year of the PVA model run cycle was estimated using the Ricker recruitment function as previously explained. The primary difference being that the value for the a parameter used in this recruitment calculation was determined from Equation 4. This determination used the $a_{\text {pop_wild }}$ and $a_{\text {pop_hatchery }}$ values assigned to each code year and the expected proportion of hatchery and wild fish. It should be noted that in most cases, the proportion of hatchery fish for the model run (expectations for future years) was different from the proportion of hatchery fish observed during the base period (code years 1972 to 1992).

The approach used to model the impact of naturally spawning hatchery fish had several key assumptions. First, when the model runs were set up, the proportion of hatchery fish into the future was fixed. The model treated this situation by adding hatchery fish to each future spawning population relative to the number of wild spawners forecast. For example, if model was set to run with a hatchery proportion of 0.33 , a forecast wild spawner escapement of 400 fish was matched with a hatchery escapement of 200 fish. If the wild escapement were 100 fish, it would be matched with a hatchery escapement of only 50 fish.

Although this approach accommodated the reproductive contribution of hatchery fish to natural production, it is an oversimplification of what most likely would occur under a real management situation. Although the number of hatchery and wild fish returning to a basin tend to share the same pattern of annual fluctuations in abundance, this relationship can be dramatically shifted if the number of hatchery smolts released is suddenly increased or decreased. Further, during periods when the natural system is producing fewer wild smolts, either because of habitat
problems or lack of escapement, the production of hatchery smolts usually remains constant. Uncorrected, this would result in a higher proportion of hatchery fish on the spawning grounds. Potentially this change would decrease the overall productivity of the population and make it more vulnerable to extinction. However, if the production of hatchery fish remains constant and a portion of them spawn in the wild, it will be theoretically impossible for the natural produced fish to disappear from the basin. As long as the hatchery program continues, there will be at least some natural recruitment. However, such a population would be entirely artificial and therefore not be consistent with the conservation of native species.

Two other issues also need qualification with respect to hatchery fish in the present form of the PVA model. Hatchery fish may be from "wildtype" broodstocks derived from local wild populations or they may be from a more "traditional-type" broodstock, typically derived from nonlocal populations and often domesticated to a certain degree. While "wild-type" hatchery fish are more likely to be genetically similar to the local wild fish than are "traditional-type" hatchery fish, there is not strong evidence that their relative reproductive capacity in the natural environment differs (Chilcote, 1998). Therefore, for the purposes of estimating natural production within the PVA model, all hatchery fish, regardless of origin, were assumed to equally capable. Information is presented later in the report that examines this assumption in more detail.

The other issue concerns what happens to the productivity of a population of mixed hatchery and wild fish when management changes are made that eliminate or greatly reduce the number of hatchery spawners. The PVA model makes adjustments in productivity under the assumption that any negative impact of hatchery fish on the overall productivity of the wild population is not permanent. It is supposed that any long-term genetic changes that have occurred in the wild population as a result of naturally spawning hatchery fish are relatively minor and will not suppress the innate productivity of the wild population.

There are several pieces of evidence that support this view. First, in those studies where it has been possible to directly measure the reproductive success of naturally spawning hatchery fish and wild fish (Chilcote et al 1986, Leider et al, 1990, and P. Hulett , personal communication), large differences between hatchery and wild fish have been found. In these particular studies, conducted on the Kalama River in Washington, wild fish retained more than a 10 -fold advantage in their productivity even though hatchery fish have been present and naturally spawning for over 20 years within the study area. Presumably if genetic damage had been taking place the accumulative effect would have
reduced that productivity of the wild spawners and the measured difference in the reproductive success between hatchery and wild fish would have been less. Consistent with this assessment are recent findings by Sharpe et al. (2000). They found biochemical evidence that wild steelhead from the Kalama River had retained a genetically distinctive identity in a comparison with the stock of hatchery fish that has been present in the Kalama basin in high numbers since the early 1970s.

Secondly, previous findings reported by Chilcote (1998) suggest that the productivity of mixed wild and hatchery populations are not particularly sensitive to the type of hatchery broodstock involved in the mix. Hatchery broodstocks from non-local, domesticated origin seem to cause a decrease in overall natural production no greater than do stocks from a local "wild-type" origin. If naturally spawning hatchery fish were causing long-term genetic damage to the wild population, it would seem that the relative greater damage from genetically dissimilar domesticated hatchery stocks versus a "wild-type" stock would be readily evident. However, this does not seem to be the case.

Finally, some hatchery programs in Oregon have been in existence for a long time and others for a relatively short time. For example, hatchery summer steelhead have been returning to the N. Umpqua since 1960 while for the Umatilla hatchery fish have been present only since 1988. If genetic damage was occurring to the wild population, it could be surmised that the longer the exposure to hatchery fish, the more adverse genetic characteristics would have accumulated. However as the results from this assessment will show, the relative productivity of mixed wild and hatchery populations does not appear to very sensitive to the length of time the mixing has taken place.

Therefore, when the available evidence is considered, it seemed reasonable to assume that once hatchery fish are removed from a natural spawning population, its productivity will increase.

The PVA model was used for several purposes, primary among these was to determine, for each population, if the criteria were met for any of the three conservation status designations, endangered, threatened, or viable. The criteria used in this report are as follows:

Endangered - Greater than $20 \%$ probability of extinction in 50 years.
Threatemed - Greater than a 5\% probability of extinction in 104 years.
Viable - Less than 5\% probability of declining below the viable abundance threshold $(0.20 / B)$ in 104 years.

It should be noted the rationale for modeling a period of 104 years instead of the more "traditional" 100-year time frame had to do with the periodicity of the survival cycle imposed by the model. To make the persistence forecasting more conservative, model runs were set up to end at the low point in the survival cycle. The 50 -year time period accomplished this. However, to end at the same point on the survival cycle for the longer model runs it was necessary to extend their length from 100 to 104 years.

The assumed fishing mortality rate for all persistence modeling was $5 \%$ for coastal populations and the Columbia basin populations downstream from Bonneville dam. For Columbia basin populations upstream of Bonneville dam the adult mortality rate was set at $15 \%$. These numbers reflect the situation that there is nearly a statewide regulation against the killing of wild steelhead. When wild fish are caught, they must be released back to the river. The $5 \%$ mortality rate assigned to these populations represents the mortality of wild fish that are caught and then die as results of handling and stress. Populations of steelhead that pass Bonneville Dam enter a non-selective gillnet fishery that causes an increased level of mortality on wild fish. Therefore, a $15 \%$ rather than a $5 \%$ mortality rate was used to model these populations.

## Assessment Results and Discussion

## Populations and Sub-populations

The data sets evaluated in this report were assembled into 31 groupings assumed to have some degree of demographic independence and therefore meeting the definition of population as proposed by McElhaney (2000). Population boundaries for Columbia Basin steelhead were comprehensively reviewed as the first step in performing this population assessment. A similar review was not done for coastal steelhead for two practical reasons. For those few locations on the coast where long-term data sets are available, the population boundaries seem relatively obvious and easy to justify as discrete, demographically independent units. The questionable population boundaries on the coast, however, are associated with basins where there is no or very little data. Because of this lack of information, doing an assessment for such populations, regardless of their assumed boundaries, was not possible. Therefore, the review of populations boundaries for such areas is lower priority and will be deferred until the necessary data becomes available in the future.

For the Columbia basin, where the data sets were more comprehensive, a total of 29 populations were identified. Six populations were identified for the Lower Columbia ESU, 6 for the Willamette ESU, 11 for the Middle Columbia, and 6 for the Snake (Appendix 1). In addition, significant sub-population structure within several of the populations was believed to exist, particularly within the Snake ESU. Population and subpopulation boundaries were based upon basin size and known discontinuities in hydrology, elevation, geology, temperature regime, vegetative cover, aspect, and spawn timing. It was assumed that these physical differences were capable of causing some degree of isolation, and genetic adaptation. In addition, a limited set of biochemical data was used to gauge the level at which sub-division among populations and sub-populations likely occurred.

Using the temporal pattern of Ricker $\boldsymbol{a}$-values calculated from moving 7year sequences of spawner-recruit data, an attempt was made to compare geographically proximate populations for evidence of demographic independence. It was assumed that one indicator of demographic independence might be dissimilar temporal patterns in population productivity. Within the Rogue basin the four populations could be classified as belonging to one of two very distinct patterns. The winter and summer steelhead populations upstream of Gold Ray dam were very similar, with a peak in productivity followed by a significant decline (Figure 1). The winter steelhead population in the Applegate and the summer steelhead population in mid Rogue followed a pattern that was nearly the inverse of the populations above Gold Ray. However, since winter and summer steelhead are assumed to be largely reproductively independent, the evidence from Rogue basin suggests that all 4 populations, either because of their life history or temporal pattern of productivity can be confidently classified as independent from each other.

The pattern for populations in the upper Willamette ESU is less clear. However, it appears that some populations like the Molalla and the Calapooia have experienced relatively wide swings in productivity over the recent past (Figure 2). In contrast, the productivity of upper South Santiam steelhead has remained relatively stable with the exception of a rapid upswing in the last several years. The lower South Santiam seems to have more similarity to the North Santiam population than it does to the steelhead in the upper South Santiam. The pattern of productivity does not support lumping the upper and lower South Santiam together as one population.

Within the Deschutes basin, the decline in productivity experienced by the Warm Springs basin steelhead in recent years was much less than
the sudden, and very large decline in productivity for steelhead monitored on the mainstem Deschutes at Sherars Falls (Figure 3). One likely, explanation of this difference is the relatively high incidence of hatchery fish in the Deschutes basin in recent years compared to the Warm Springs system where only wild fish are allowed into the basin.

Six populations were identified for the John Day basin. The temporal pattern of productivity among these populations appears strikingly similar during the middle portion of the observed time series, with the exception of steelhead from the upper North Fork John Day (Figure 4). This population did not show the same decline experienced by populations in the rest of the basin. Because of this difference, it seems reasonable to treat this group of fish as a population separate from those in the nearby lower North Fork or Middle Fork John Day. The productivity of steelhead in the Middle Fork, South Fork and upper mainstem seemed to track quite closely, while steelhead from the lower mainstem and the lower North Fork were different in their productivity during the beginning of the time series. From the standpoint of similar trends in productivity and geographically proximity, if any of these populations should be lumped it would be upper mainstem and the South Fork. However, arguing against such a recommendation is biochemical evidence that suggests that steelhead in the South Fork are genetically distinct within the John Day basin (Kostow, 1995). Therefore, it was concluded that the boundaries for the populations in the John Day should remain as they are described in this report (Appendix 1).

Although 4 populations were provisionally identified within the Grande Ronde basin, data were available from only 3 locations. The productivity of steelhead in the middle mainstem section (Phillips Creek index site) had quite high productivity during the first part to the time series however, by the end of the time series had declined to nearly the same low productivity as steelhead from the upper Grande Ronde (Figure 5). Although the Joseph population experienced a moderate decline towards the end of the time series, the inception of this decline seemed to lag several years behind productivity decline observed for the other the Grande Ronde populations. The magnitude and pattern of productivity among these three groups of steelhead appears as great, if not greater, than for groups of fish that were classified as separate populations in the John Day basin. Therefore, it was concluded the steelhead from the middle Grande Ronde section should not be lumped into the same population as those from the upper Grand Ronde. The geographically logical alternative would be to assign the middle mainstem group to the lower Grande Ronde population. This would result in the moving of the lower Grande Ronde population upstream to encompass the Lookingglass and middle mainstem sub-populations.

## Population Trends

The trend in annual pre-harvest abundance of wild fish was examined for 31 populations. In some cases, such as the Hood River and Walla Walla populations, the time series was too short for meaningful evaluation. However, for most populations it was possible to look at the pattern of wild fish abundance for the last 20 to 30 years. Nearly all populations had a rapid decline in abundance during the early to mid 1990s and a low point in abundance during the late 1990s (Appendix 2). However, beyond this shared characteristic there appeared to be 3 semi-distinct temporal patterns of steelhead abundance. By far the most common pattern (Type 1) is characterized by a period of low abundance, followed by a period of greater abundance, and then most recently a second, but more severe low period. To varying degrees all of the populations in the Middle Columbia appear to display this pattern. This pattern also appears to be weakly displayed by several of the coastal populations including the Salmonberry and the upper Rogue summer-run. The second pattern observed (Type 2) seems to be predominating in the Snake ESU. The Type 2 pattern is similar to the Type 1 , however in the case of the Type 2 the first period of low abundance is deeper than the second low abundance period. A third pattern (Type 3) was also recognized. It was characterized by a steady decline with no peak in abundance or evidence of cyclic character. This pattern appears most commonly for steelhead populations in the Upper Willamette and Lower Columbia ESUs. Finally 3 populations did not appear to follow any of these patterns. Two populations, winter steelhead in the North Umpqua and upper Rogue displayed a similar cyclic abundance pattern, with no overall trend up or down (Appendix 2). They appeared relatively stable. The Applegate population appeared unique in that it remained relatively stable during the early 1990 s, unlike nearly every other population that showed a declined during this period. In addition, the overall trend for the Applegate population appears to upward.

## Observed Abundance and Conservation Thresholds

As described in the methods section, critical and viable abundance thresholds were determined for those populations with sufficient abundance data to perform a PVA analyses. Twenty-seven populations met this requirement. In comparing the observed average abundance of wild fish for these populations over the last 6 years, all were greater than the thresholds for critical and viable (Table 1). Indeed, 7 populations appeared to be at abundance levels greater than needed for maximum
production of recruits, (i.e., maximum seeding). These included 4 coastal populations (Rogue winter-run, Applegate, N. Umpqua summer-run, and N. Umpqua winter-run), one from the mid-Columbia, the upper N. Fork John Day, and 2 Snake basin populations, Joseph and the lower Grande Ronde. Of the remaining populations, 9 had abundance numbers in the range of $100 \%$ to $50 \%$ of maximum seeding, while 11 were in the range between $50 \%$ of maximum seeding and the viable threshold.

Table 1. Observed 6 -year average wild steelhead abundance and conservation abundance thresholds, for 27 populations of wild steelhead in Oregon.
Abundance expressed as either total spawners (data without decimal points) or as spawners per stream mile (data with decimal points).

| Population | Maximum <br> seeding | so\% <br> Seeding | Viable <br> Threshold | Critical <br> Threshold | Recent <br> 6-yr <br> Average |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Upper Rogue SR | 4485 | 2242 | 897 | 275 | 3142 |
| Upper Rogue WR | 4343 | 2172 | 869 | 247 | 7352 |
| Mid Rogue SR | 47.1 | 23.5 | 9.4 | 6.3 | 17.6 |
| Applegate WR | 1048 | 524 | 210 | 63 | 1371 |
| N. Umpqua SR | 3233 | 1617 | 647 | 189 | 3546 |
| N. Umpqua WR | 4273 | 2137 | 855 | 234 | 6692 |
| Salmonberry | 7.2 | 3.6 | 1.4 | 0.5 | 4.8 |
| Sandy | 1677 | 839 | 336 | 82 | 651 |
| Clackamas | 1396 | 698 | 279 | 73 | 395 |
| Molalla | 49.8 | 24.9 | 9.9 | 2.6 | 14.0 |
| N. Santiam | 83.9 | 41.5 | 16.6 | 13.0 | 21.9 |
| Lower S. Santiam | 41.3 | 20.6 | 8.1 | 2.1 | 8.4 |
| Upper S. Santiam | 524 | 262 | 108 | 33 | 312 |
| Calapooia | 11.3 | 5.6 | 2.2 | 0.8 | 8.3 |
| Deschutes | 7394 | 3697 | 1149 | 398 | 1997 |
| Warm Springs | 399 | 199 | 80 | 32 | 162 |
| Lower John Day | 3.9 | 2.0 | 0.79 | 0.2 | 2.68 |
| Lower NFk John Day | 4.3 | 2.2 | 0.9 | 0.2 | 2.62 |
| Upper NFk John Day | 2.3 | 1.1 | 0.5 | 0.1 | 3.0 |
| Middle Fork John Day | 11.2 | 5.6 | 2.2 | 0.8 | 4.8 |
| South Fork John Day | 8.4 | 4.2 | 1.7 | 0.5 | 2.6 |
| Upper John Day | 7.7 | 3.9 | 1.5 | 0.5 | 2.6 |
| Umatilla | 1666 | 833 | 333 | 103 | 1247 |
| Joseph | 3.4 | 1.7 | 0.7 | 0.2 | 4.6 |
| Lower Grande Ronde | 2.2 | 1.1 | 0.8 | 0.3 | 2.2 |
| Upper Grande Ronde | 3.9 | 1.9 | 0.5 | 0.1 | 3.3 |
| Imnaha | 6.0 | 3.0 | 1.2 | 0.4 | 4.7 |

## Trends in Productivity

A temporal series of Ricker a parameter estimates were obtained for each population (see methods section). These are presented for all populations in Appendix 3. It was assumed that the pattern of these $\boldsymbol{a}$ parameters related directly to the pattern of productivity and recruitment for each population over time. Although there were considerable differences among populations (Figures 1, 2, 3, 4 and 5), the overall pattern suggested the existence of cyclic phenomenon with respect to recruitment. As shown in Figure 6, it appeared that a negative slope cycle, with period of approximately 18 years, explained much of the observed annual variation in the recruitment performance of these populations. As might be expected, this pattern in recruitment is similar to the pattern of pre-harvest abundance for many populations.

Although annual variations in the freshwater environment have been shown to influence the long-term recruitment pattern in steelhead in northern British Columbia (Smith, 2000), it is unlikely that this is the primary controlling factor for Oregon populations. Ecological fluctuations in the marine environment that effect steelhead survival seems to be the best explanation. The observed cyclic pattern of productivity holds for Oregon steelhead from a wide range of geographically distinct and variable freshwater habitats. If variations in the freshwater environment were the primary controlling factor, the heterogeneous nature of these habitats across Oregon would not lead to a nearly universal pattern of fluctuating productivity. Smith and Ward (2000) and Welch et al. (2000) both present evidence that marine survival conditions can be a major factor in controlling the recruitment of steelhead.

Given the apparent importance of this single factor, it is clear that population assessments that do not incorporate the natural cyclic variability in the quality of the marine environment for steelhead will yield potentially erroneous results. This was one of the reasons the PVA model used in this assessment was structured around an assumed cyclic pattern of marine survivals. The sequence of Ricker a parameters estimated for the multiple 7-year data sets for each population was used as a means to approximate this cyclic phenomenon.

## Population Viability Analysis

In examining the PVA results, only the Deschutes and North Santiam populations were found to have probabilities of extinction high enough to trigger the criteria for the classification for endangered (Table 2). Four of the 27 populations, including the North Santiam and Deschutes, met the
criteria for threatened. These same 4 populations (Middle Rogue summer-run, N. Santiam, Deschutes, and Umatilla) did not meet the criteria for viable. All had a 5\% or greater probability that at the end of the next 104 years their abundance level would be less than the viable threshold $(0.20 / \boldsymbol{B})$. However, the remaining 22 populations examined all appear at relatively low risk and have greater than $95 \%$ chance of remaining above their respective viable threshold levels. It should be emphasized that these findings are based upon two key assumptions concerning the future productivity of the 27 populations modeled.

Table 2. The probability of extinction and the probability of an abundance less than the viable threshold for 27 populations of steelhead in Oregon with respect to criteria for the classification of endangered, threatened, and viable as determined from PVA modeling.

| Population | Viable | Threatened | Endangered |
| :--- | :---: | :---: | :---: |
| Upper Rogue SR | 0.00 | 0.00 | 0.00 |
| Upper Rogue WR | 0.00 | 0.00 | 0.00 |
| Mid Rogue SR | $\mathbf{0 . 2 8}$ | $\mathbf{0 . 1 9}$ | 0.00 |
| Applegate WR | 0.00 | 0.00 | 0.00 |
| N. Umpqua SR | 0.00 | 0.00 | 0.00 |
| N. Umpqua WR | 0.00 | 0.00 | 0.00 |
| Salmonberry | 0.00 | 0.00 | 0.00 |
| Sandy | 0.00 | 0.00 | 0.00 |
| Clackamas | 0.00 | 0.00 | 0.00 |
| Molalla | 0.00 | 0.00 | 0.00 |
| N. Santiam | $\mathbf{0 . 7 8}$ | $\mathbf{0 . 6 3}$ | 0.13 |
| Lower S. Santiam | 0.00 | 0.00 | 0.00 |
| Upper S. Santiam | 0.00 | 0.00 | 0.00 |
| Calapooia | 0.00 | 0.00 | 0.00 |
| Deschutes | $\mathbf{1 . 0 0}$ | $\mathbf{0 . 9 9}$ | $\mathbf{0 . 8 5}$ |
| Warm Springs | 0.01 | 0.00 | 0.00 |
| Lower John Day | 0.00 | 0.00 | 0.00 |
| Lower NFk John Day | 0.00 | 0.00 | 0.00 |
| Upper NFk John Day | 0.00 | 0.00 | 0.00 |
| Middle Fork John Day | 0.00 | 0.00 | 0.00 |
| South Fork John Day | 0.00 | 0.00 | 0.00 |
| Upper John Day | 0.00 | 0.00 | 0.00 |
| Umatilla | $\mathbf{1 . 0 0}$ | $\mathbf{0 . 9 8}$ | $\mathbf{0 . 5 4}$ |
| Joseph | 0.00 | 0.00 | 0.00 |
| Lower Grande Ronde | 0.00 | 0.00 | 0.00 |
| Upper Grande Ronde | 0.00 | 0.00 | 0.00 |
| Imnaha | 0.00 | 0.00 | 0.00 |

First, it was assumed the survival pattern in future years would be cyclic and be no less than $30 \%$ of the values observed for individual populations from 1973 to 1995, the baseline period.

Secondly, it was assumed changes in the proportion of hatchery fish in the naturally spawning population would have an effect on the future productivity of the population. This component was included in the model runs because the supposed relationship between the proportion of naturally spawning hatchery fish and overall population productivity was confirmed by the data (see Hatchery Impacts section). However, as described in this later section, the direct or indirect impact of hatchery fish on natural production was found to be much greater than expected. How this difference was added to the model runs was critical to outcomes for populations that involved hatchery fish. In particular, model runs for the Deschutes and Umatilla populations were extremely sensitive to how much reproductive discounting was applied to naturally spawning hatchery fish. Regardless, using the standard discounting approach described in the methods section, the results of supplemental PVA model runs suggested that if the future proportion of naturally spawning fish in the Deschutes and Umatilla populations was reduced by approximately $1 / 3$, the probability of extinction would decrease to less than 0.05 .

## Assessment Synthesis

None of the populations examined meet the numerical abundance thresholds for threatened or endangered. In addition, the PVA model results suggest that only four of these populations are at significant conservation risk, populations in the middle Rogue, North Santiam, Deschutes, and Umatilla.

In the early 1990s, most populations entered a period of decline. For populations in the lower Columbia and upper Willamette ESUs, this decline appears to have been a feature that started prior to 1990. However, the record for the majority of other populations in Oregon, provides evidence that this decline may be part of a normal cyclic pattern. Rather than a chronic, long-term decline, as appears the case for the Willamette and lower Columbia populations, the pattern observed for most other populations suggests a long-term cyclic phenomena. Indeed, in the last 5 years several populations appear to be entering the ascending portion of this cycle.

The greatest concentration of vulnerable populations appeared to be those that belonged to the mid-Columbia ESU. Two populations, the

Deschutes and Umatilla, met the criteria for an endangered classification. A majority of the populations in this ESU are at abundance levels that are less than $50 \%$ of maximum seeding. Nearly equal, in terms of vulnerability, were the Upper Willamette populations. Only did 2 out 5 of these populations were at levels of escapement greater than necessary for $50 \%$ of maximum seeding. In addition, one population, the North Santiam, met the criteria for a threatened classification. Although, the PVA analysis did not suggest that the two populations representing the lower Columbia ESU, the Sandy and Clackamas, were at risk of extinction, these populations show other troubling signs. Both exhibit a chronic downward trend in abundance with little indication an underlying cyclic pattern exists that might reverse this trend. In addition, within the last 6 years, both populations have experienced at least one escapement of wild fish that was less than the viable threshold. Therefore, these populations may be more vulnerable than the PVA analysis seems to suggest.

In a less vulnerable category than the populations discussed in the previous paragraph, are those belonging to the KMP, the Oregon Coast and the Snake ESUs. The KMP is represented by 4 populations within the Rogue basin. Two of the populations, both winter-runs, appear quite healthy and are currently at levels greater than maximum seeding. The summer steelhead population upstream of Gold Ray Dam, while in considerably greater abundance than the level necessary for $50 \%$ of maximum seeding, has experienced a drop in numbers and now appear to be stabilizing at a new, lower level. The density of summer steelhead downstream of Gold Ray Dam is much lower than historical levels. They met the criteria for threatened classification. However, in recent years, particularly 2000, there has been a substantial increase over the extremely low spawner densities observed in the early 1990s.

This primary problem for the Oregon Coast ESU is that long-term data sets exist for only 3 populations, and 2 of these are found in the same basin, the Umpqua. Regardless, the winter and summer steelhead populations in the North Umpqua appear healthy. For both populations, the current number of wild fish exceeds the level necessary for maximum production. However, the only population north of the Umpqua for which there is adequate data, is the Salmonberry. Although, this population has a 6 -year abundance average that is more the level necessary for $50 \%$ of maximum seeding, record low spawner densities have been experienced the last 3 years of this time series. While several ODFW biologists have expressed concern that these low values may be an artifact of modified stream survey methodologies, a means for accommodating these changes has not yet been devised.

All of the populations examined within the Snake ESU appear to be at abundance levels that are greater than $50 \%$ of maximum seeding. Both the Joseph and Imnaha populations have survived a period of extremely low spawner densities in the late 1970s. They are now substantially above these levels and seem to be in the beginning stages of an upward trend. The pattern for the other 2 Grande Ronde populations is more erratic. The upper Grande Ronde spawner density in the last 2 years has been very low. However, the productivity for these populations has remained greater than for many other populations during the recent low portion of the presumed survival cycle (Figures 1 through 5).

In terms of their conservation status, steelhead populations appear to fall into one of two groups. The healthier of the two groups contains steelhead populations belonging to the Snake, Oregon Coast, and KMP ESUs. Although not without problems and areas of concern, they are not at risk of extinction. Less healthy, is the group that contains populations belonging to the Middle Columbia, Lower Columbia, and Upper Willamette ESUs. Several of these populations are at substantial risk of extinction.

## Mortality Rate Assessment

To assess the impact of human-caused fish mortality (e.g., fisheries) on the status and recovery of steelhead within the Columbia basin, a series of PVA model runs were performed for a range of different assumed adult mortality rates. For each population the probability of extinction over a 50 -year time period was estimated for 16 mortality rates between $0 \%$ and $75 \%$. In performing these model runs it should be clarified that the human mortality being discussed is primarily related to fisheries. Since this evaluation is based upon spawner-recruit relationships of the last 20 years, it incorporates a certain degree of background mortality associated with human activities such as the operation of dams and other adverse land uses. However, the mortality from these other activities is not directly accounted in this modeling exercise. Such accounting would be possible if reliable estimates for past and future levels of these additional human-caused mortalities were available. However, in this analysis, the future magnitude of these other sources of mortality was assumed to be the same as what they had been in the past.

For mortality rates between $0 \%$ and $20 \%$, the probability of extinction remained at 0.00 for nearly all populations except the Middle Rogue, N . Santiam, Deschutes, and Umatilla populations (Table 3). The of the four exceptions, the Deschutes population appeared most vulnerable to mortalities imposed by fisheries. However, the actual vulnerability is probably less than these results suggest; particularly if this reduction in
mortality rate was made by closing all steelhead fishing in the Deschutes basin．Such a closure would have its largest impact on the number of hatchery spawners in the basin．Under current regulations，hatchery steelhead may be caught and kept．However，all wild steelhead caught must be released．A consequence of closing this fishery would be to increase the proportion of naturally spawning hatchery fish in the basin， which would further reduce the population＇s productivity．Therefore，it is possible that the small gain in reduced survival could be cancelled by a substantial decrease in population productivity．

Table 3．PVA simulations of estimated probability of extinction in 50 years for 27 populations of Oregon steelhead under 16 different hypothetical adult mortality rates．

| Population | Percent Adult Mortality Rate |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0 | 5 | 10 | 15 | 20 | 25 | 30 | 35 | 40 | 45 | 50 | 55 | 60 | 65 | 70 | 75 |
| Rogue SR |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 02 | ， 90 | 140 |  |
| MidRogueSR | ． 00 | 01 | 11 | 41 | 88 | 99 | 14 | － 7 d | \％${ }^{\text {dex }}$ | 6804 | 4 | 4， | \％ 6 | 1 | W |  |
| Rogue WR |  |  |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 84 |  |
| Applegate |  |  |  |  |  |  |  |  | ． 00 | $\stackrel{46}{ }$ | \％ 6 | 4䋂 | 5098 | \％ 6 | － |  |
| NUmp SR |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 43 | 新新 | \％${ }^{4}$ |  |
| Nump WR |  |  |  |  |  |  |  |  |  |  |  |  | ． 00 | ． 04 | 99 | 484 |
| Salmonberry |  |  |  |  |  |  |  | ． 00 | 04 | .25 | 73 | 540 | 40 | \％ 14 | － 8 | \％ 6 |
| Calapooia |  |  |  | ． 00 | 01 | ． 08 | 12 | 30 | 59 | 88 | 98 | ${ }^{18}$ | \％ 6 | \％${ }^{\text {a }}$ | 1 | 468 |
| LoS．Santiam |  |  |  |  |  |  |  |  | ． 00 | 02 | 03 | ． 05 | 29 | 51 | 75 | ． 92 |
| UpS．Santiam |  |  |  |  |  |  |  |  |  | ． 00 | 05 | 51 | ． 95 | 94 |  | 46\％ |
| N．Santiam | ． 03 | .11 | 31 | 63 | 88 | 99 | 30 | 4 | \％ | 318 | 619 | 31974 | 4， 0 | \％ 4 | \％－9 | \％${ }^{2}$ |
| Molalla |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 01 | 20 | 68 | ． 88 |
| Clackamas |  |  |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 10 | ． 91 |
| Sandy |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 01 | 20 | 72 | ¢ |
| WarmSprings |  |  |  |  | ． 00 | ． 01 | 09 | 28 | 77 | 99 | \％ | 569 | W0． | Y， 6 | － |  |
| Deschutes | .36 | 53 | 69 | 85 | .91 | 98 | 99 | 0 | 10 | 4， | 4 4 | 10 | \％11 | 10 | 20］ |  |
| LowerNF JohnDay |  |  |  |  |  |  |  |  |  | ． 00 | 01 | ． 34 | 84 | 10 | 1020 | 16 |
| Upper NF John Day |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ． 00 |
| M．Fork John Day |  |  |  |  |  | ． 00 | 02 | 14 | ． 38 | 77 | 99 | W | 448 | 109 | 49 | 464 |
| S．Fork John Day |  |  |  |  |  |  | ． 00 | 05 | 17 | ． 53 | 92 | 914 | 40 | H2\％ | 40 | －${ }^{\text {¢ }}$ |
| LowerMainsJohnDay |  |  |  |  |  |  | ． 00 | ． 01 | 04 | 10 | 13 | 27 | 47 | 69 | 94 | 10 |
| UprMainsJohn Day |  |  |  |  | ． 00 | 01 | ． 04 | 18 | .41 | 62 | 90 | 99 | 109 | 56 | 948 | 4， |
| Umatilla | 01 | 08 | 21 | 57 | 85 | 98 | 1.0 | 10 | 1.0 | 4 4 | 296\％ | 4－9 | 4 | 10 | －${ }^{18}$ | 10 |
| UprGrRond |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 03 | 27 | 78 | 99 |
| MidGrRond |  |  |  |  |  |  | ． 00 | ． 01 | 13 | 77 | 2918 | \％ 9 | \％ 6 | 51 | Y 8 | 14 |
| Joseph |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ． 00 | 29 |
| Imnaha |  |  |  |  |  |  |  |  |  | ． 00 | 01 | 09 | 74 | Y 9 | 19 | － |

Therefore，closing a sport fishery in order to gain，in most cases，no more than a $5 \%$ reduction in wild fish mortality，would be counter productive in basins where hatchery fish are commonly caught and removed from the population．In such basins，such a strategy would likely make the status of the wild population worse instead of better．The interpretation
of the results presented in Table 3 should keep this practical management consideration in mind.

For most populations, the model predicted the risk of extinction was very low until the mortality rates reached $30 \%$. However, once the mortality rate became greater than $40 \%$, the probability of extinction among the remaining populations began to increase. There was considerable difference among populations at what mortality rate the probability of extinction started to increase.

There also appeared to be a threshold mortality rate at which the transition to a $100 \%$ probability of extinction was rapid. For most populations, once a mortality rate was found that increased the probability of extinction above 0.00, an increment of an additional 15\% to $25 \%$ in mortality rate was usually sufficient to result in a probability of extinction of 1.00 . For example, the probability of extinction for the Warm Springs population under a $25 \%$ mortality rate was 0.01 . However, the probability of extinction increased to 0.99 when the model was run using a $45 \%$ mortality rate. Therefore, it appears that while most of these populations can withstand moderate levels of adult mortality with no significant impact on their likelihood of persistence, once the mortality level goes past the trigger point, the probability of extinction increases suddenly.

Since the transition from low risk to high risk happens so rapidly once the critical mortality rate is exceeded, management strategies should set a limit on maximum mortality rates at some level considerable less than this trigger point. To do otherwise leaves no room for logic errors in the model used to forecast these impacts, nor does it allow for any error in the actual measurement of mortality rates.

For most populations the trigger point is a mortality rate of $30 \%$ or higher. Taking a conservative approach, it seems that a mortality rate limit of $20 \%$ is a reasonable conservation standard for most steelhead populations in Oregon.

The mortality rate assessment described here was based upon adult mortalities. Therefore, it would seem best suited for evaluating the impact of various fishery management strategies. However, various sampling, monitoring, and collection activities will also impact steelhead at the juvenile life history stage. One way to account for the impact of mortality on juvenile steelhead would be to convert each life history stage (i.e., eggs, fry, parr, and smolts) into adult equivalents. Once stated in terms of adult equivalents they could be converted into a cumulative mortality rate and a determination made if the population was under the critical management level (e.g., 20\%).

However, there are some serious flaws with this approach. First, for the majority of populations in the Columbia basin the method of population assessment is redd surveys. For a variety of technical reasons, converting these redd per stream mile index counts into a spawner estimate for an entire population with any reliability is very difficult, at least with the current state of the knowledge about this species and the habitat characteristics of each basin. Second, without a reliable total run-size or spawner estimate, juvenile mortalities converted into adult equivalents have limited value because it would not be possible to estimate the proportion they represent of the total population. Third, converting to adult equivalents means assigning an expected survival rate for each juvenile life history stage to adulthood. Although such survival rates have been estimated, they are highly variable for many reasons including: spawner densities, habitat quality, climatic variations, inter and intra-specific competition, predation, and disease.

A more viable approach for assessing the impact of activities that cause juvenile mortality is to base them on a direct estimate of the mortality rate, rather than numbers in adult equivalents. For example, the number of smolts killed in the course of operating a smolt trap can be expressed as a mortality rate. Or if 1 out of every 50 fry encountered during electroshocking surveys are killed then this can be converted to a mortality rate. In the latter example, this mortality rate would be weighed in proportion to the amount of steelhead habitat for the entire population. However, even if this was not possible, localized estimates could be used as the maximum likely mortality rate for the entire population, a very conservative, yet feasible approach.

Although, the survival rates of fry and parr to the smolt stage are higher when the spawner seeding level is lower (density dependent survival), one could make the conservative assumption that this survival is density independent at all life history stages. Essentially, this is the assumption that the habitat is extremely underseeded and there is no competition for space and food among juvenile steelhead. With this assumption, the life history stage at which mortality occurs is not a complication as long as it is expressed in terms of a mortality rate for the entire population. Under this assumption of density independent survival, if $10 \%$ of the members of a population are killed because of some management action, the impact is the same, regardless if it happens at the fry, smolt, or adult stages.

With this operating assumption, the mortality across all life history stages could be combined into one cumulative mortality rate using a simple equation such as:

Cumulative mortality rate $=1-\left[\left(1-\mathrm{M}_{\mathrm{f}}\right)^{*}\left(1-\mathrm{M}_{\mathrm{p}}\right)^{*}\left(1-\mathrm{M}_{\mathrm{s}}\right)^{*}\left(1-\mathrm{M}_{\mathrm{a}}\right)\right]$
Where $\mathrm{M}_{\mathrm{f}}, \mathrm{M}_{\mathrm{p}}, \mathrm{M}_{\mathrm{s}}$, and $\mathrm{M}_{\mathrm{a}}$ equal the estimated mortality rate at the life history stage of fry, parr, smolt, and adult, respectively. Using this approach, once the cumulative, life history mortality rate was calculated for a population, the resulting probability of high risk could be estimated directly from population specific PVA modeling results such as those shown in Table 3. For example, a cumulative mortality rate of $30 \%$ for the North Santiam population corresponds with an extinction probability of 0.25 , exceeding the criteria for an endangered classification (i.e., a 0.20 probability of extinction. Therefore, if a cumulative mortality rate of $30 \%$ occurred for the North Santiam population, its conservation status could be expected to change from sensitive to endangered.

## Hatchery Impacts

For many of the populations assessed, hatchery fish are present in the production areas used by wild fish and spawn naturally. In comparing such mixed populations, it appears that the higher the proportion of hatchery fish, the poorer the subsequent recruitment of naturally produced offspring (Chilcote, 1998). However, from 1998 to present several major changes in hatchery programs were made to reduce the number of naturally spawning hatchery fish. Some of these changes are expected to eliminate naturally spawning hatchery fish altogether, especially for populations in the Lower Columbia and Upper Willamette. Therefore, to model the future status of such populations with the PVA methodology, it was necessary to make a positive adjustment in their reproductive potential to account for the removal of the hatchery fish. The procedure for accomplishing this has been previously described in the Analytical Concepts and Methods section of this report. The discussion that follows here describes some of the key findings that were a byproduct of developing this adjustment procedure for the PVA modeling.

The relationship between the proportion of hatchery fish in 15 natural populations and their respective productivity was evaluated for 12 different time intervals (Table 4). This comparison was based on 15 populations, 12 of which had hatchery fish present on the spawning grounds at some point during the last 25 years. In addition, data had been collected on each of these 12 populations to measure proportion of hatchery fish. The remaining 3 populations were comprised only of wild fish. They were selected to provide some reference points for the "no hatchery fish" condition.

In all 12 cases, a negative relationship was found; the higher the proportion of hatchery fish in the spawning population, the more population productivity declined, as measured by the value for the a parameter in the Ricker recruitment function (Figures 7 through 9). Nine of the 12 regressions were statistically significant (Table 5). Of the 3 that were not significant, the common problem was an anomalous data point for the Sandy population (Figures 7 and 8). For the 3 time intervals in question, it was decided that the regressions without the Sandy data would be used. The rationale for this decision was based upon a closer examination of the data points from which the a parameters were estimated for the Sandy population between the brood years 1982 and 1990. This period was characterized by a rapid transition to very low productivity and rather large escapements. This combination, by chance, yielded a very steep regression line which intersected the y axis (the y-intercept is the estimate for at an unlikely high value. When the Sandy data point was removed from these 3 regressions, they all became statistically significant.

It should also be noted that not all regressions were based upon the data from 15 populations. For the earlier time periods, this was because the data set for the Molalla and the N. Santiam did not start until the 1980 and 1983 brood years, respectively (Table 4). For the later time periods, fewer populations were represented because for some populations with an older age structure, the total number of recruits for the 1994 and 1995 brood years can not be estimated until that adult returns for 2001 or 2002 have been counted (Table 4).

The results presented here lead to the conclusion that overall population productivity can be adversely effected by naturally spawning hatchery fish. Further that this effect is not minor. For nearly all of the time intervals evaluated, it appears that when the proportion of hatchery fish exceeds $60 \%$, the population can no longer replace its self, even at very low densities where the recruitment function would predict that survival would be at its greatest.

It is unclear whether the mechanism for this relationship is genetic or environmental. However, if it is genetic, the use of wild fish from local populations for hatchery broodstock does not appear to be a corrective solution. The evidence for this statement is in the distribution of data points around the regression lines describing the negative relationship between productivity and hatchery fish proportion for each of the 12 time intervals evaluated (Figures 7 through 9). Some of these data points represent populations with hatchery fish that were derived from local, wild fish. In contrast, other data points are from populations where the hatchery fish are of non-local origin and may be partially domesticated.

Table 4. Steelhead populations used in comparing productivity and proportion of hatchery fish over 12 time intervals, "x" denotes inclusion of population in regression analysis.

| Population | Time Interval for Regressions |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & 78 \\ & \text { to } \\ & 84 \end{aligned}$ | $\begin{aligned} & 79 \\ & \text { to } \\ & 85 \end{aligned}$ | $\begin{aligned} & 80 \\ & \text { to } \\ & 86 \end{aligned}$ | $\begin{aligned} & \hline 81 \\ & \text { to } \\ & 87 \end{aligned}$ | $\begin{array}{\|l} \hline 82 \\ \text { to } \\ 88 \end{array}$ | $\begin{array}{\|l} \hline 83 \\ \text { to } \\ 89 \end{array}$ | $\begin{aligned} & 84 \\ & \text { to } \\ & 90 \end{aligned}$ | $\begin{aligned} & 85 \\ & \text { to } \\ & 91 \end{aligned}$ | $\begin{aligned} & \hline 86 \\ & \text { to } \\ & 92 \end{aligned}$ | $\begin{aligned} & 87 \\ & \text { to } \\ & 93 \end{aligned}$ | $\begin{aligned} & 88 \\ & \text { to } \\ & 94 \end{aligned}$ | $\begin{aligned} & \hline 89 \\ & \text { to } \\ & 95 \end{aligned}$ |
| Joseph | x | x | x | x | x | x | x | x | x | x |  |  |
| Imnaha | x | x | x | x | x | x | x | x | x | x | x |  |
| Upr Grande Ronde | x | x | X | X | x | x | x | x | x | X |  |  |
| Lwr NFk. John Day | x | x | x | x | x | x | x | x | x | x | x |  |
| Umatilla | x | x | x | x | x | x | x | x | x | x | x |  |
| Deschutes | x | x | x | x | x | x | x | x | x | x |  |  |
| Sandy | x | x | x | x | x | x | x | x | x | x | x | x |
| Clackamas | x | x | x | x | x | x | x | x | x | X | x | x |
| Molalla |  |  | x | x | x | x | x | x | x | x | x | x |
| North Santiam |  |  |  |  |  | x | x | x | x | x | x | x |
| Upr S. Santiam | x | x | x | x | x | x | x | x | x | x | x | x |
| N.Umpqua SR | x | x | x | x | x | x | x | x | x | x | x | x |
| N.Umpqua WR | x | x | x | x | x | x | x | x | x | x | x |  |
| Rogue SR | x | x | x | X | x | X | x | x | x | x | x | x |
| Rogue WR | x | x | X | x | x | x | X | x | x | x | x | x |

Table 5. Statistics for linear regressions of population productivity and proportion of naturally spawning hatchery fish based upon 15 Oregon steelhead populations over 12 time intervals.

| Time Interval | $\mathbf{n}$ | $\mathbf{R}^{\mathbf{2}}$ | $\mathbf{p}$ | Intercept | Slope |
| :--- | :---: | :---: | :---: | :---: | :---: |
| $1978-84$ | 13 | 0.39 | 0.022 | 2.130 | -3.700 |
| $1979-85$ | 13 | 0.74 | 0.000 | 2.180 | -4.270 |
| $1980-86$ | 14 | 0.76 | 0.000 | 2.040 | -3.270 |
| $1981-87$ | 14 | 0.72 | 0.000 | 2.080 | -3.060 |
| $\mathbf{1 9 8 2 - 8 8} \mathbf{w /} /$ Sandy | $\mathbf{1 4}$ | $\mathbf{0 . 1 5}$ | $\mathbf{0 . 1 7 5}$ | $\mathbf{1 . 8 6 8 1}$ | $\mathbf{- 1 . 3 7 7}$ |
| $1982-88 \mathrm{w} / \mathrm{o}$ Sandy | 13 | 0.70 | 0.000 | 1.920 | -2.870 |
| $\mathbf{1 9 8 3 - 8 9 \mathrm { w } / \text { Sandy }}$ | $\mathbf{1 5}$ | $\mathbf{0 . 1 6}$ | $\mathbf{0 . 1 2 7}$ | $\mathbf{1 . 4 9 0 1}$ | $\mathbf{- 2 . 2 8 3}$ |
| $1983-89 \mathrm{w} / \mathrm{o}$ Sandy | 14 | 0.56 | 0.002 | 1.620 | -4.440 |
| $\mathbf{1 9 8 4 - 9 0} \mathbf{~} /$ Sandy | $\mathbf{1 5}$ | $\mathbf{0 . 1 9}$ | $\mathbf{0 . 0 9 6}$ | $\mathbf{1 . 2 4 1 2}$ | $\mathbf{- 2 . 8 3 2 9}$ |
| $1984-90 \mathrm{w} / \mathrm{o}$ Sandy | 14 | 0.62 | 0.001 | 1.420 | -5.270 |
| $1985-91$ | 15 | 0.47 | 0.005 | 0.770 | -3.420 |
| $1986-92$ | 15 | 0.55 | 0.002 | 1.010 | -4.480 |
| $1987-93$ | 15 | 0.66 | 0.001 | 1.050 | -3.320 |
| $1988-94$ | 11 | 0.70 | 0.000 | 1.192 | -3.898 |
| $1989-95$ | 8 | 0.93 | 0.000 | 1.705 | -4.106 |

If the use of wild fish for hatchery broodstock is a strategy that improves the genetic adaptation and reproductive success of naturally spawning hatchery fish, then the data points from these types of programs should consistently appear above the regression line (more productive than the average). The corollary would be that the data points from populations that contain hatchery fish from non-local, domesticated sources, should consistently fall below the regression line (less productive than the average).

However, this is not the pattern that was observed. In the regression analyses performed, the populations with hatchery fish that have been predominately non-local origin are the upper Grand Ronde, Deschutes, Sandy, Clackamas, and Molalla. Those populations with local, "wildtype" hatchery fish are the Imnaha, Umatilla, N. Santiam, S. Santiam, Umpqua summer-run, Rogue summer-run and Rogue winter-run. If the data points from all 12 regressions (time intervals) are combined, 39 points can be assigned to populations with non-local hatchery fish and 63 to populations with "wild-type" hatchery stocks. When these points were further classified as to whether they fall above or below the regression line, the two types of populations did not differ. For populations with "wild-type" hatchery fish, $60 \%$ of the data points were above the regression line. For populations with non-local, domesticated hatchery fish, $62 \%$ of the data points were above the regression line.

These results lead to one of two conclusions. Either that the use of wild fish in hatchery programs does solve the genetic problem that makes hatchery fish genetically maladapted for natural survival. A conclusion that implies rapid and significant genetic change occurs when fish are brought into the hatchery environment.

Alternatively, that the use of wild fish for hatchery broodstock greatly reduces the genetic difference between hatchery and wild fish, but this really doesn't matter because the mechanism causing the reduced productivity for naturally spawning hatchery is not genetic. The problem is caused by some unknown environmental impact of the hatchery rearing environment that results in hatchery fish being less able to produce viable offspring under natural conditions.

Regardless of the mechanism, when hatchery fish mix with wild fish in natural production areas, the overall productivity of the population declines. In effect the freshwater habitat becomes less efficient in producing steelhead. Not only does this mean that natural production goals are compromised, it means that the population's vulnerability to extinction is increased.

## Zones of Inference

It was not possible to perform an assessment on every steelhead population in Oregon. This was either due to lack of representative data or data sets that did cover enough years for the analytical approach used here. In particular, specific coverage was not possible for much of the Oregon coast as well as portions of the Grande Ronde and Imnaha basins.

However, in examining the results for populations for which there were data, there was not a great deal of variation with respect their status within each ESU. With certain exceptions, such as the Deschutes, the consistency of the assessment results suggests that the zone of inference concerning the biological health for these populations is probably at the ESU level. Therefore, where specific information on a specific population or sub-population does not exist, it is reasonable to infer that its status is probably similar to that of other populations within the same ESU for which an assessment exists.

In addition, the sensitivity of steelhead populations to mortality (fishery or other sources) appears relatively consistent, again with a few exceptions. Therefore, in terms of mortality impact, the zone of inference is sufficiently broad to conclude that as long as the mortality rate does not exceed $20 \%$, the probability of extinction is very low (the model results suggest zero).

Finally, with respect to hatchery programs, the impact of naturally spawning hatchery fish on the capacity of a population to produce recruits appears universally adverse. Therefore, the zone of inference concerning the impact of naturally spawning hatchery on wild populations is statewide. Without specific data to the contrary, it is a reasonable inference that wild steelhead populations are better off when returning hatchery fish are prevented from escaping into natural spawning areas.

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Figure 1. Temporal distribution of Ricker $a$-values estimated from 7brood year sequences of spawner recruit data from 1980 to 96 for 4 populations of Rogue River steelhead.


Figure 2. Temporal distribution of Ricker $\mathfrak{a}$-values estimated from 7brood year sequences of spawner recruit data from 1980 to 96 for 5 populations of Willamette River steelhead.


Figure 3. Temporal distribution of Ricker a-values estimated from 7brood year sequences of spawner recruit data from 1980 to 96 for 5 populations of Deschutes River steelhead.


Figure 4. Temporal distribution of Ricker od-values estimated from 7brood year sequences of spawner recruit data from 1980 to 96 for 6 populations of John Day River steelhead.


Figure 5. Temporal distribution of Ricker $\mathfrak{a}$-values estimated from 7brood year sequences of spawner recruit data from 1980 to 96 for 3 populations of Grande Ronde River steelhead.


Figure 6. Temporal distribution of Ricker a-values estimated from 7brood year sequences of spawner-recruit data from 1972 to 1993 brood cycle code years for 15 populations of Oregon steelhead comprised of less than $15 \%$ hatchery fish.







Figure 7. Relationship between productivity (vertical axis) and proportion of hatchery fish in spawning population (horizontal axis) for 15 steelhead populations for 5 time intervals. Last two panels of graph from 1982-88 time interval, with and without data point for Sandy population.


Figure 8. Relationship between productivity (vertical axis) and proportion of hatchery fish in spawning population (horizontal axis) for 15 steelhead populations for 5 time intervals. First 4 panels of graph represent data with and without Sandy population for the 1982-88 and 1983-89 time intervals.


Figure 9. Relationship between productivity (vertical axis) and proportion of hatchery fish in spawning population (horizontal axis) for 15 steelhead populations for 5 time intervals. First 4 panels of graph represent data with and without Sandy population for the 1982-88 and 1983-89 time intervals.

Appendix 1. Presumed steelhead populations and sub-populations in Oregon's portion of the Columbia River basin.

| ESU | Population | Subpopulation | Description |
| :---: | :---: | :---: | :---: |
| Upper Willamette | Calapooia |  |  |
|  | Lower S. Santiam |  | Basin from N. Santiam to Wiley Creek (downstream from Foster Dam) |
|  | Upper S.Santiam |  | Basin from Foster Dam upstream |
|  | North Santiam |  |  |
|  | West Valley | Luckiamute |  |
|  |  | Rickreall |  |
|  |  | Yamhill |  |
|  |  | Tualatin |  |
|  | Molalla |  |  |
|  |  | + | + |
| Lower Columbia | Scappose |  |  |
|  | Clackamas | Abernathy |  |
|  |  | Clackamas |  |
|  | Sandy | Sandy |  |
|  | Columbia Gorge | Tanner |  |
|  |  | Eagle |  |
|  |  | Herman |  |
|  |  | Lindsey |  |
|  | Hood WR |  | Winter Steelhead |
|  | Hood SR |  | Summer Steelhead |
|  |  |  | Were |
| Middle Columbia | Fifteenmile | Chenowith | Winter Steelhead |
|  |  | Fifteenmile | Winter Steelhead |
|  | Deschutes |  | Basin except Warm Springs |
|  | Warm Springs |  |  |
|  | Lower John Day |  | Basin from mouth to South Fork John Day, exclusive of North Fork John Day |
|  | Lower North Fork John Day |  | North Fork Basin from Mainstem John Day to Middle Fork John Day |
|  | Upper North Fork John Day |  | North Fork Basin from Middle Fork John Day upstream |
|  | Middle Fork John Day |  |  |
|  | South Fork John Day |  |  |
|  | Upper John Day |  | John Day Basin upstream of South Fork John Day |
|  | Umatilla | Lower Umatilla | Basin downstream McKay Creek |
|  |  | Upper Umatilla | Basin upstream from McKay Creek except Meacham Creek drainage |


|  |  | Meacham Creek |  |
| :---: | :---: | :---: | :---: |
|  | Walla Walla | Middle Walla Walla | Basin above Touchet R. up and including Cottonwood Creek |
|  |  | Upper Walla Walla | Basin above Cottonwood Creek |
|  |  | 4 | - |
| Snake | Lower Grande Ronde | Wenaha |  |
|  |  | Lower Mainstem | Grande Ronde basin from state line to (and including) Phillips Crk, exclusive of Wenaha, Joseph Crk, and Lookingglass Crk |
|  |  | Lookingglass |  |
|  | Joseph |  |  |
|  | Wallowa | North Wallowa | Wallowa tributaries on northern side of basin up to Prairie Creek |
|  |  | South Wallowa | Wallowa tributaries on southern side of basin, exclusive of Minam |
|  |  | Prairie |  |
|  |  | Minam |  |
|  | Upper Grande Ronde | Middle Mainstem | Grande Ronde básin tributaries from Phillips Creek to upper end of Grande Ronde valley (near the city of La Grande) exclusive of Catherine and Willow Creeks |
|  |  | Willow |  |
|  |  | Catherine |  |
|  |  | Upper Mainstem | Grande Ronde basin tributaries from upper end of Grande Ronde valley (near the city of La Grande) up to and including Meadow Creek. |
|  |  | South Upper Mainstem | Grande Ronde basin upstream from Meadow Creek. |
|  | Imnaha | Zumwalt | Camp Creek and tributaries on the west side of the basin from downstream from Big Sheep Creek. |
|  |  | Lower Imnaha | Tributaries on the eastern portion of the basin downstream from Big Sheep Creek. |
|  |  | Big Sheep |  |
|  |  | Upper Imnaha | Basin upstream of Big Sheep Creek |
|  | Snake | Hell's Canyon | Oregon tributaries from state line to Hell's Canyon Dam. |

Appendix 2. "Populations at a glance" information summaries for 27 populations of steelhead within Oregon.

| Basin: Imnaha |
| :---: |
| Population: Imnaha |
| Sub-population: Zumwalt |
| Monitoring sites: Camp Creek |
|  |
| Method: Redd Surveys |


Average Distribution of Ages in return year

Repeat Age 2 Age 3 Age 4 Age 5 Age 6 | Females per Redd | $=0.81$ |
| ---: | :--- |
| Prop. of females in spng pop. | $=0.60$ |

| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre-Harv Wild Fish | 6 -yr. <br> Moving Av |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Spnrs |  | Columbia | In-basin | Combined |  |  |
| 1974 | 2.30 | 0.00 | 3.11 | 0.00 | 3.11 | 0.289 | 0.02 | 0.30 | 4.5 | 2.4 |
| 1975 | 0.70 | 0.00 | 0.95 | 0.00 | 0.95 | 0.128 | 0.02 | 0.15 | 1.1 | 1.9 |
| 1976 | 0.20 | 0.00 | 0.27 | 0.00 | 0.27 | 0.067 | 0.02 | 0.09 | 0.3 | 2.0 |
| 1977 | 1.00 | 0.00 | 1.35 | 0.00 | 1.35 | 0.078 | 0.02 | 0.10 | 1.5 | 1.8 |
| 1978 | 1.80 | 0.00 | 2.43 | 0.00 | 2.43 | 0.208 | 0.02 | 0.22 | 3.1 | 2.1 |
| 1979 | 2.70 | 0.00 | 3.65 | 0.00 | 3.65 | 0.196 | 0.02 | 0.21 | 4.6 | 2.5 |
| 1980 | 5.70 | 0.00 | 7.70 | 0.00 | 7.70 | 0.079 | 0.02 | 0.10 | 8.5 | 3.2 |
| 1981 | 1.50 | 0.00 | 2.03 | 0.00 | 2.03 | 0.087 | 0.02 | 0.11 | 2.3 | 3.4 |
| 1982 | 1.20 | 0.00 | 1.62 | 0.00 | 1.62 | 0.069 | 0.02 | 0.09 | 1.8 | 3.6 |
| 1983 | 2.80 | 0.00 | 3.78 | 0.00 | 3.78 | 0.069 | 0.02 | 0.09 | 4.1 | 4.1 |
| 1984 | 2.30 | 0.00 | 3.11 | 0.00 | 3.11 | 0.088 | 0.02 | 0.11 | 3.5 | 4.1 |
| 1985 | 6.50 | 0.00 | 8.78 | 0.00 | 8.78 | 0.121 | 0.02 | 0.14 | 10.2 | 5.1 |
| 1986 | 7.20 | 0.00 | 9.72 | 0.00 | 9.72 | 0.209 | 0.02 | 0.23 | 12.5 | 5.7 |
| 1987 | 8.56 | 2.14 | 11.56 | 2.89 | 14.45 | 0.139 | 0.02 | 0.16 | 13.7 | 7.6 |
| 1988 | 13.44 | 3.36 | 18.14 | 4.54 | 22.68 | 0.158 | 0.02 | 0.17 | 22.0 | 11.0 |
| 1989 | 6.56 | 1.64 | 8.86 | 2.21 | 11.07 | 0.172 | 0.02 | 0.19 | 10.9 | 12.1 |
| 1990 | 10.40 | 2.60 | 14.04 | 3.51 | 17.55 | 0.161 | 0.02 | 0.18 | 17.1 | 14.4 |
| 1991 | 3.04 | 0.76 | 4.10 | 1.03 | 5.13 | 0.160 | 0.02 | 0.18 | 5.0 | 13.5 |
| 1992 | 1.44 | 0.36 | 1.94 | 0.49 | 2.43 | 0.147 | 0.02 | 0.16 | 2.3 | 11.8 |
| 1993 | 5.44 | 1.36 | 7.34 | 1.84 | 9.18 | 0.164 | 0.02 | 0.18 | 9.0 | 11.0 |
| 1994 | 5.28 | 1.32 | 7.13 | 1.78 | 8.91 | 0.155 | 0.02 | 0.17 | 8.6 | 8.8 |
| 1995 | 1.76 | 0.44 | 2.38 | 0.59 | 2.97 | 0.105 | 0.02 | 0.12 | 2.7 | 7.4 |
| 1996 | 2.24 | 0.56 | 3.02 | 0.76 | 3.78 | 0.106 | 0.02 | 0.12 | 3.5 | 5.2 |
| 1997 | 2.72 | 0.68 | 3.67 | 0.92 | 4.59 | 0.090 | 0.02 | 0.11 | 4.1 | 5.0 |
| 1998 | 4.32 | 1.08 | 5.83 | 1.46 | 7.29 | 0.105 | 0.02 | 0.12 | 6.6 | 5.8 |
| 1999 | 1.60 | 0.40 | 2.16 | 0.54 | 2.70 | 0.090 | 0.02 | 0.11 | 2.4 | 4.7 |
| 2000 | 5.92 | 1.48 | 7.99 | 2.00 | 9.99 | 0.079 | 0.02 | 0.10 | 8.9 | 4.7 |


| Basin: Grande Ronde |
| :---: |
| Population: Joseph |
| Sub-population: |
| Monitoring sites: Butte Creek |
| Crow Creek |
| Elk Creek |
| Peavine Creek |
| Swamp Creek |
| Method: Redd Surveys |



|  | Critical Threshold |
| :--- | :--- |
| Viable Threshold | 0.19 |
|  | 0.67 |
|  | 4.59 |


| Females per Redd | $=0.81$ |
| ---: | :--- |


| Spawning <br> Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre- Hanv e-Hanv Wild F | 6 year |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwrrs | Hatch Spnrs |  | Columbia | 1 n -basin | Combined |  |  |
| 1974 | 1.6 | 0.0 | 2.1 | 0.0 | 2.1 | 0.289 | 0.02 | 0.30 | 3.0 |  |
| 1975 | 0.5 | 0.0 | 0.6 | 0.0 | 0.6 | 0.128 | 0.02 | 0.15 | 0.7 |  |
| 1976 | 0.5 | 0.0 | 0.6 | 0.0 | 0.6 | 0.067 | 0.02 | 0.09 | 0.7 |  |
| 1977 | 0.9 | 0.0 | 1.2 | 0.0 | 1.2 | 0.078 | 0.02 | 0.10 | 1.3 |  |
| 1978 | 0.7 | 0.0 | 1.0 | 0.0 | 1.0 | 0.208 | 0.02 | 0.22 | 1.3 |  |
| 1979 | 0.2 | 0.0 | 0.3 | 0.0 | 0.3 | 0.196 | 0.02 | 0.21 | 0.4 | 1.2 |
| 1980 | 2.6 | 0.0 | 3.6 | 0.0 | 3.6 | 0.079 | 0.02 | 0.10 | 3.9 | 1.4 |
| 1981 | 1.8 | 0.0 | 2.4 | 0.0 | 2.4 | 0.087 | 0.02 | 0.11 | 2.7 | 1.7 |
| 1982 | 2.0 | 0.0 | 2.7 | 0.0 | 2.7 | 0.069 | 0.02 | 0.09 | 3.0 | 2.1 |
| 1983 | 1.2 | 0.0 | 1.6 | 0.0 | 1.6 | 0.069 | 0.02 | 0.09 | 1.8 | 2.2 |
| 1984 | 1.8 | 0.0 | 2.4 | 0.0 | 2.4 | 0.088 | 0.02 | 0.11 | 2.7 | 2.4 |
| 1985 | 7.6 | 0.0 | 10.3 | 0.0 | 10.3 | 0.121 | 0.02 | 0.14 | 11.9 | 4.3 |
| 1986 | 7.6 | 0.0 | 10.3 | 0.0 | 10.3 | 0.209 | 0.02 | 0.23 | 13.3 | 5.9 |
| 1987 | 6.3 | 0.0 | 8.6 | 0.0 | 8.6 | 0.139 | 0.02 | 0.16 | 10.1 | 7.1 |
| 1988 | 7.8 | 0.0 | 10.5 | 0.0 | 10.5 | 0.158 | 0.02 | 0.17 | 12.7 | 8.7 |
| 1989 | 7.5 | 0.0 | 10.2 | 0.0 | 10.2 | 0.172 | 0.02 | 0.19 | 12.5 | 10.5 |
| 1990 | 7.0 | 0.0 | 9.5 | 0.0 | 9.5 | 0.161 | 0.02 | 0.18 | 11.5 | 12.0 |
| 1991 | 1.2 | 0.0 | 1.6 | 0.0 | 1.6 | 0.160 | 0.02 | 0.18 | 1.9 | 10.3 |
| 1992 | 2.1 | 0.0 | 2.9 | 0.0 | 2.9 | 0.147 | 0.02 | 0.16 | 3.4 | 8.7 |
| 1993 | 5.5 | 0.0 | 7.5 | 0.0 | 7.5 | 0.164 | 0.02 | 0.18 | 9.1 | 8.5 |
| 1994 | 2.1 | 0.0 | 2.8 | 0.0 | 2.8 | 0.155 | 0.02 | 0.17 | 3.4 | 7.0 |
| 1995 | 1.8 | 0.0 | 2.4 | 0.0 | 2.4 | 0.105 | 0.02 | 0.12 | 2.7 | 5.4 |
| 1996 | 1.6 | 0.0 | 2.2 | 0.0 | 2.2 | 0.106 | 0.02 | 0.12 | 2.5 | 3.9 |
| 1997 | 2.3 | 0.0 | 3.1 | 0.0 | 3.1 | 0.090 | 0.02 | 0.11 | 3.5 | 4.1 |
| 1998 | 4.7 | 0.0 | 6.4 | 0.0 | 6.4 | 0.105 | 0.02 | 0.12 | 7.3 | 4.8 |
| 1999 | 3.8 | 0.0 | 5.2 | 0.0 | 5.2 | 0.090 | 0.02 | 0.11 | 5.8 | 4.2 |
| 2000 | 3.82 | 0.0 | 5.2 | 0.0 | 5.2 | 0.079 | 0.02 | 0.10 | 5.7 | 4.6 |


| Basin: Grande Ronde |
| :---: |
| Population: Lower Grande Ronde |
| Sub-population: Middle Mainstem |
| Monitoring sites: Phillips Creek |
|  |
| Method: Redd Surveys |


Prop. of females in sprg pop. $=0.60$

| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre- Harv Wild Fish | $\begin{gathered} \text { 6-yT} \\ \text { Moving Av } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Sprrs |  | Columbia | In-basin | Combined |  |  |
| 1967 | 5.60 | 0.00 | 7.56 | 0.00 | 7.56 | 0.21 | 0.02 | 0.23 | 9.8 |  |
| 1968 | 1.20 | 0.00 | 1.62 | 0.00 | 1.62 | 0.21 | 0.02 | 0.23 | 2.1 |  |
| 1969 | 4.40 | 0.00 | 5.94 | 0.00 | 5.94 | 0.21 | 0.02 | 0.23 | 7.7 |  |
| 1970 | 2.80 | 0.00 | 3.78 | 0.00 | 3.78 | 0.21 | 0.02 | 0.23 | 4.9 |  |
| 1971 | 2.40 | 0.00 | 3.24 | 0.00 | 3.24 | 0.257 | 0.02 | 0.27 | 4.4 |  |
| 1972 | 1.20 | 0.00 | 1.62 | 0.00 | 1.62 | 0.227 | 0.02 | 0.24 | 2.1 |  |
| 1973 | 4.40 | 0.00 | 5.94 | 0.00 | 5.94 | 0.301 | 0.02 | 0.32 | 8.7 | 5.0 |
| 1974 | 0.80 | 0.00 | 1.08 | 0.00 | 1.08 | 0.289 | 0.02 | 0.30 | 1.5 | 4.9 |
| 1975 | 1.60 | 0.00 | 2.16 | 0.00 | 2.16 | 0.128 | 0.02 | 0.15 | 2.5 | 4.0 |
| 1976 | 1.20 | 0.00 | 1.62 | 0.00 | 1.62 | 0.067 | 0.02 | 0.09 | 1.8 | 3.5 |
| 1977 | 2.00 | 0.00 | 2.70 | 0.00 | 2.70 | 0.078 | 0.02 | 0.10 | 3.0 | 3.3 |
| 1978 | 4.00 | 0.00 | 5.40 | 0.00 | 5.40 | 0.208 | 0.02 | 0.22 | 7.0 | 4.1 |
| 1979 | 0.40 | 0.00 | 0.54 | 0.00 | 0.54 | 0.196 | 0.02 | 0.21 | 0.7 | 2.7 |
| 1980 | 5.60 | 0.00 | 7.56 | 0.00 | 7.56 | 0.079 | 0.02 | 0.10 | 8.4 | 3.9 |
| 1981 | 1.20 | 0.00 | 1.62 | 0.00 | 1.62 | 0.087 | 0.02 | 0.11 | 1.8 | 3.8 |
| 1982 | 0.80 | 0.00 | 1.08 | 0.00 | 1.08 | 0.069 | 0.02 | 0.09 | 1.2 | 3.7 |
| 1983 | 0.80 | 0.00 | 1.08 | 0.00 | 1.08 | 0.069 | 0.02 | 0.09 | 1.2 | 3.4 |
| 1984 | 2.56 | 0.00 | 3.46 | 0.00 | 3.46 | 0.088 | 0.02 | 0.11 | 3.9 | 2.9 |
| 1985 | 2.40 | 0.00 | 3.24 | 0.00 | 3.24 | 0.121 | 0.02 | 0.14 | 3.8 | 3.4 |
| 1986 | 4.80 | 0.00 | 6.48 | 0.00 | 6.48 | 0.209 | 0.02 | 0.23 | 8.4 | 3.4 |
| 1987 | 6.00 | 0.00 | 8.10 | 0.00 | 8.10 | 0.139 | 0.02 | 0.16 | 9.6 | 4.7 |
| 1988 | 1.70 | 0.30 | 2.30 | 0.41 | 2.70 | 0.158 | 0.02 | 0.17 | 2.8 | 4.9 |
| 1989 | 1.36 | 0.24 | 1.84 | 0.32 | 2.16 | 0.172 | 0.02 | 0.19 | 2.3 | 5.1 |
| 1990 | 1.70 | 0.30 | 2.30 | 0.41 | 2.70 | 0.161 | 0.02 | 0.18 | 2.8 | 4.9 |
| 1991 | 2.48 | 0.44 | 3.35 | 0.59 | 3.94 | 0.160 | 0.02 | 0.18 | 4.1 | 5.0 |
| 1992 | 5.44 | 0.96 | 7.34 | 1.30 | 8.64 | 0.147 | 0.02 | 0.16 | 8.8 | 5.0 |
| 1993 | 1.36 | 0.24 | 1.84 | 0.32 | 2.16 | 0.164 | 0.02 | 0.18 | 2.2 | 3.8 |
| 1994 | 1.02 | 0.18 | 1.38 | 0.24 | 1.62 | 0.155 | 0.02 | 0.17 | 1.7 | 3.6 |
| 1995 | 1.70 | 0.30 | 2.30 | 0.41 | 2.70 | 0.105 | 0.02 | 0.12 | 2.6 | 3.7 |
| 1996 | 1.36 | 0.24 | 1.84 | 0.32 | 2.16 | 0.106 | 0.02 | 0.12 | 2.1 | 3.6 |
| 1997 | 0.68 | 0.12 | 0.92 | 0.16 | 1.08 | 0.090 | 0.02 | 0.11 | 1.0 | 3.1 |
| 1998 | 2.72 | 0.48 | 3.67 | 0.65 | 4.32 | 0.105 | 0.02 | 0.12 | 4.2 | 2.3 |
| 1999 | 1.36 | 0.24 | 1.84 | 0.32 | 2.16 | 0.090 | 0.02 | 0.11 | 2.1 | 2.3 |
| 2000 | 0.68 | 0.12 | 0.92 | 0.16 | 1.08 | 0.079 | 0.02 | 0.10 | 1.0 | 2.2 |


| Basin: Grande Ronde |
| :---: |
| Population: Upper G. Ronde |
| Sub-population: Upper Mainstem |
| Monitoring sites: McCoy Creek |
| Meadows Greek |
| Method: Redd Surveys |




|  | Average Distribution of Ages in return year |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
|  | 0.03 | 0.00 | 0.02 | 0.38 | 0.44 | 0.13 |
| Mile | Effective | Harvest Rates |  |  | Pre-Hañ | 6 year |
| Hatch Spnrs | Spawners | Columbia | In-basin | Combined | Wild Fish | Moving Av. |
| 0.00 | 15.96 | 0.21 | 0.02 | 0.23 | 20.6 |  |


| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre- Hanv Wild Fish | 6 year <br> Moving Av. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Spnrs |  | Columbia | In-basin | Combined |  |  |
| 1967 | 11.8 | 0.0 | 15.96 | 0.00 | 15.96 | 0.21 | 0.02 | 0.23 | 20.6 |  |
| 1968 | 1.9 | 0.0 | 2.61 | 0.00 | 2.61 | 0.21 | 0.02 | 0.23 | 3.4 |  |
| 1969 | 4.3 | 0.0 | 5.74 | 0.00 | 5.74 | 0.21 | 0.02 | 0.23 | 7.4 |  |
| 1970 | 2.3 | 0.0 | 3.08 | 0.00 | 3.08 | 0.21 | 0.02 | 0.23 | 4.0 |  |
| 1971 | 4.3 | 0.0 | 5.83 | 0.00 | 5.83 | 0.257 | 0.02 | 0.27 | 8.0 |  |
| 1972 | 4.9 | 0.0 | 6.61 | 0.00 | 6.61 | 0.227 | 0.02 | 0.24 | 8.7 |  |
| 1973 | 5.4 | 0.0 | 7.23 | 0.00 | 7.23 | 0.301 | 0.02 | 0.32 | 10.6 | 7.0 |
| 1974 | 0.4 | 0.0 | 0.58 | 0.00 | 0.58 | 0.289 | 0.02 | 0.30 | 0.8 | 6.6 |
| 1975 | 1.1 | 0.0 | 1.49 | 0.00 | 1.49 | 0.128 | 0.02 | 0.15 | 1.7 | 5.6 |
| 1976 | 0.3 | 0.0 | 0.38 | 0.00 | 0.38 | 0.067 | 0.02 | 0.09 | 0.4 | 5.0 |
| 1977 | 0.7 | 0.0 | 0.92 | 0.00 | 0.92 | 0.078 | 0.02 | 0.10 | 1.0 | 3.9 |
| 1978 | 1.0 | 0.0 | 1.35 | 0.00 | 1.35 | 0.208 | 0.02 | 0.22 | 1.7 | 2.7 |
| 1979 | 0.3 | 0.0 | 0.34 | 0.00 | 0.34 | 0.196 | 0.02 | 0.21 | 0.4 | 1.0 |
| 1980 | 2.2 | 0.0 | 2.94 | 0.00 | 2.94 | 0.079 | 0.02 | 0.10 | 3.3 | 1.4 |
| 1981 | 0.1 | 0.0 | 0.07 | 0.00 | 0.07 | 0.087 | 0.02 | 0.11 | 0.1 | 1.2 |
| 1982 | 0.1 | 0.0 | 0.09 | 0.00 | 0.09 | 0.069 | 0.02 | 0.09 | 0.1 | 1.1 |
| 1983 | 1.8 | 0.0 | 2.46 | 0.00 | 2.46 | 0.069 | 0.02 | 0.09 | 2.7 | 1.4 |
| 1984 | 2.8 | 0.0 | 3.83 | 0.00 | 3.83 | 0.088 | 0.02 | 0.11 | 4.3 | 1.8 |
| 1985 | 12.5 | 0.0 | 16.93 | 0.00 | 16.93 | 0.121 | 0.02 | 0.14 | 19.6 | 5.0 |
| 1986 | 2.0 | 0.0 | 2.75 | 0.00 | 2.75 | 0.209 | 0.02 | 0.23 | 3.5 | 5.1 |
| 1987 | 4.0 | 0.0 | 5.40 | 0.00 | 5.40 | 0.139 | 0.02 | 0.16 | 6.4 | 6.1 |
| 1988 | 3.7 | 0.5 | 4.97 | 0.68 | 5.64 | 0.158 | 0.02 | 0.17 | 6.8 | 7.2 |
| 1989 | 1.2 | 0.2 | 1.57 | 0.21 | 1.78 | 0.172 | 0.02 | 0.19 | 2.2 | 7.2 |
| 1990 | 1.6 | 0.2 | 2.16 | 0.29 | 2.46 | 0.161 | 0.02 | 0.18 | 3.0 | 6.9 |
| 1991 | 1.7 | 0.5 | 2.31 | 0.69 | 2.99 | 0.160 | 0.02 | 0.18 | 3.6 | 4.3 |
| 1992 | 1.7 | 0.5 | 2.30 | 0.69 | 2.99 | 0.147 | 0.02 | 0.16 | 3.6 | 4.3 |
| 1993 | 1.8 | 0.6 | 2.49 | 0.74 | 3.23 | 0.164 | 0.02 | 0.18 | 3.9 | 3.9 |
| 1994 | 0.7 | 0.2 | 0.97 | 0.29 | 1.26 | 0.155 | 0.02 | 0.17 | 1.5 | 3.0 |
| 1995 | 1.1 | 0.3 | 1.45 | 0.43 | 1.88 | 0.105 | 0.02 | 0.12 | 2.1 | 3.0 |
| 1996 | 1.5 | 0.4 | 1.97 | 0.59 | 2.56 | 0.106 | 0.02 | 0.12 | 2.9 | 3.0 |
| 1997 | 3.0 | 0.9 | 4.01 | 1.20 | 5.20 | 0.090 | 0.02 | 0.11 | 5.8 | 3.3 |
| 1998 | 3.6 | 1.1 | 4.86 | 1.45 | 6.32 | 0.105 | 0.02 | 0.12 | 7.2 | 3.9 |
| 1999 | 0.6 | 0.2 | 0.78 | 0.23 | 1.01 | 0.090 | 0.02 | 0.11 | 1.1 | 3.5 |
| 2000 | 0.24 | 0.07 | 0.32 | 0.10 | 0.42 | 0.079 | 0.02 | 0.10 | 0.5 | 3.3 |


| Basin: Walla Walla |
| :---: |
| Population: Walla Walla |
| Sub-population: |
| Monitoring sites: Trap @ Nursery |
| Bridge |
| Method: Direct counts and |
| mark-recapture |
| estimates |




|  | Average Distribution of Ages at time of spawning |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |  |  |
|  | 0.05 | 0.00 | 0.29 | 0.48 | 0.18 | 0.00 |  |  |
|  |  |  |  |  |  |  |  | Wild Abund 6-yт <br> Moving Avg |
| Spawning |  |  | Effective | Wild Fish Harvest Rates |  |  | Pre- Harv |  |
| Year | SpwnrsWild | SpwnrsHatc | Tot. Spwnrs | Out-basin | In-basin | Combined | e-Han Fish Fi |  |
| 1993 | 815 | 2 | 817 | 0.16 | 0.010 | 0.17 | 985 |  |
| 1994 | 535 | 1 | 536 | 0.16 | 0.010 | 0.16 | 640 |  |
| 1995 | 430 | 5 | 435 | 0.11 | 0.010 | 0.11 | 485 |  |
| 1996 | 358 | 7 | 365 | 0.11 | 0.010 | 0.11 | 404 |  |
| 1997 | 292 | 5 | 297 | 0.09 | 0.010 | 0.10 | 324 |  |
| 1998 | 378 | 3 | 381 | 0.10 | 0.010 | 0.11 | 426 |  |
| 1999 | 279 | 1 | 280 | 0.09 | 0.010 | 0.10 | 310 | 432 |
| 2000 | 514 | 13 | 527 | 0.08 | 0.010 | 0.09 | 564 | 419 |


| Basin: Umatilla |
| :---: |
| Population: Umatilla |
| Sub-population: |
| Monitoring sites: Threemile Dam |
|  |
|  |
| Trap |
| Method: Total count of |
| returning fish. |


| Critical Threshold | 140 |
| :--- | :--- |
|  | Viable Threshold |
| Last 6-yr Average | 333 |
|  | 1247 |


| Spawning | Average Distribution of Ages at time of spawning |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
|  |  |  | 0.05 | 0.00 | 0.29 | 0.48 | 0.18 | 0.00 |
|  | Note: Spawner numbers account for wild and hatchery fish removed for broodstock at 3-mile dam - as does pre-harv abundance |  |  |  |  |  |  |  |
|  |  |  | Effective | Wild F | sh Harves | Rates | Pre- Harv | 6 -yr |
|  | SpwnrsWild | SpwrrsHatc | Tot. Spwnrs | Out-basin | In-basin | Combined | -Harv Wild Fi | Moving Avg |
| 1973 | 2057 | - | 2057 | 0.301 | 0.12 | 0.39 | 3346 |  |
| 1974 | 2640 | 0 | 2640 | 0.289 | 0.12 | 0.37 | 4217 |  |
| 1975 | 2171 | 0 | 2171 | 0.128 | 0.12 | 0.23 | 2830 |  |
| 1976 | 2534 | 0 | 2534 | 0.067 | 0.12 | 0.18 | 3086 |  |
| 1977 | 1258 | 0 | 1258 | 0.078 | 0.12 | 0.19 | 1551 |  |
| 1978 | 3080 | 0 | 3080 | 0.208 | 0.12 | 0.30 | 4421 | 3242 |
| 1979 | 2337 | 0 | 2337 | 0.196 | 0.12 | 0.29 | 3304 | 3235 |
| 1980 | 2367 | 0 | 2367 | 0.079 | 0.12 | 0.19 | 2919 | 3019 |
| 1981 | 1218 | 0 | 1218 | 0.087 | 0.12 | 0.20 | 1516 | 2800 |
| 1982 | 608 | 0 | 608 | 0.069 | 0.12 | 0.18 | 742 | 2409 |
| 1983 | 1103 | 0 | 1103 | 0.069 | 0.12 | 0.18 | 1346 | 2375 |
| 1984 | 2262 | 0 | 2262 | 0.088 | 0.12 | 0.20 | 2819 | 2108 |
| 1985 | 3093 | 0 | 3093 | 0.121 | 0.12 | 0.23 | 3998 | 2223 |
| 1986 | 2816 | 0 | 2816 | 0.209 | 0.12 | 0.30 | 4047 | 2411 |
| 1987 | 3296 | 0 | 3296 | 0.139 | 0.12 | 0.24 | 4348 | 2883 |
| 1988 | 2183 | 166 | 2349 | 0.158 | 0.12 | 0.26 | 2946 | 3251 |
| 1989 | 1944 | 371 | 2315 | 0.172 | 0.12 | 0.27 | 2668 | 3471 |
| 1990 | 1315 | 246 | 1561 | 0.161 | 0.12 | 0.26 | 1781 | 3298 |
| 1991 | 625 | 387 | 1012 | 0.160 | 0.01 | 0.17 | 751 | 2757 |
| 1992 | 2010 | 523 | 2533 | 0.147 | 0.01 | 0.16 | 2381 | 2479 |
| 1993 | 1172 | 616 | 1788 | 0.164 | 0.01 | 0.17 | 1417 | 1991 |
| 1994 | 853 | 345 | 1198 | 0.155 | 0.01 | 0.16 | 1020 | 1669 |
| 1995 | 789 | 656 | 1445 | 0.105 | 0.01 | 0.11 | 890 | 1373 |
| 1996 | 1196 | 785 | 1981 | 0.106 | 0.01 | 0.11 | 1351 | 1302 |
| 1997 | 906 | 1463 | 2369 | 0.090 | 0.01 | 0.10 | 1006 | 1344 |
| 1998 | 773 | 802 | 1575 | 0.105 | 0.01 | 0.11 | 872 | 1093 |
| 1999 | 1024 | 661 | 1685 | 0.090 | 0.01 | 0.10 | 1136 | 1046 |
| 2000 | 2032 | 713 | 2745 | 0.079 | 0.01 | 0.09 | 2229 | 1247 |


| Basin: John Day |
| :---: |
| Population: Lower NF John Day |
| Sub-population: |
| Monitoring sites: Wall Cr |
| Wilson Cr |
|  |
| Method: Redd Surveys |


|  | Critical Threshold |
| :--- | :--- |
| Viable Threshold | 0.24 |
|  | 0.86 |
|  | $\mathbf{2 . 9 0}$ |


| Females per Redd | $=0.81$ |
| ---: | :--- |
| Prop. of females in spng pop. | $=0.60$ |


Average Distribution of Ages in return year

| Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.05 | 0.00 | 0.41 | 0.43 | 0.11 | 0.00 |


| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre- Harv Abundance | 6 year Moving Av |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Spris |  | Columbia | John Day | Combined |  |  |
| 1976 | 4.75 | 0.00 | 6.4 | 0.0 | 6.4 | 0.067 | 0.12 | 0.18 | 7.8 |  |
| 1977 | 4.50 | 0.00 | 6.1 | 0.0 | 6.1 | 0.078 | 0.12 | 0.19 | 7.5 |  |
| 1978 | 5.09 | 0.00 | 6.9 | 0.0 | 6.9 | 0.208 | 0.12 | 0.30 | 9.9 |  |
| 1979 | 0.12 | 0.00 | 0.2 | 0.0 | 0.2 | 0.196 | 0.12 | 0.29 | 0.2 |  |
| 1980 | 6.81 | 0.00 | 9.2 | 0.0 | 9.2 | 0.079 | 0.12 | 0.19 | 11.3 |  |
| 1981 | 4.81 | 0.00 | 6.5 | 0.0 | 6.5 | 0.087 | 0.12 | 0.20 | 8.1 | 7.5 |
| 1982 | 4.06 | 0.00 | 5.5 | 0.0 | 5.5 | 0.069 | 0.12 | 0.18 | 6.7 | 7.3 |
| 1983 | 1.41 | 0.00 | 1.9 | 0.0 | 1.9 | 0.069 | 0.12 | 0.18 | 2.3 | 6.4 |
| 1984 | 1.76 | 0.00 | 2.4 | 0.0 | 2.4 | 0.088 | 0.12 | 0.20 | 3.0 | 5.3 |
| 1985 | 13.90 | 0.00 | 18.8 | 0.0 | 18.8 | 0.121 | 0.12 | 0.23 | 24.3 | 9.3 |
| 1986 | 12.11 | 0.00 | 16.4 | 0.0 | 16.4 | 0.209 | 0.12 | 0.30 | 23.5 | 11.3 |
| 1987 | 12.14 | 0.00 | 16.4 | 0.0 | 16.4 | 0.139 | 0.12 | 0.24 | 21.6 | 13.6 |
| 1988 | 2.87 | 0.00 | 3.9 | 0.0 | 3.9 | 0.158 | 0.12 | 0.26 | 5.2 | 13.3 |
| 1989 | 1.11 | 0.00 | 1.5 | 0.0 | 1.5 | 0.172 | 0.12 | 0.27 | 2.1 | 13.3 |
| 1990 | 0.71 | 0.00 | 1.0 | 0.0 | 1.0 | 0.161 | 0.12 | 0.26 | 1.3 | 13.0 |
| 1991 | 1.36 | 0.00 | 1.8 | 0.0 | 1.8 | 0.160 | 0.04 | 0.19 | 2.3 | 9.3 |
| 1992 | 2.29 | 0.00 | 3.1 | 0.0 | 3.1 | 0.147 | 0.12 | 0.25 | 4.1 | 6.1 |
| 1993 | 0.27 | 0.00 | 0.4 | 0.0 | 0.4 | 0.164 | 0.12 | 0.26 | 0.5 | 2.6 |
| 1994 | 1.85 | 0.00 | 2.5 | 0.0 | 2.5 | 0.155 | 0.04 | 0.19 | 3.1 | 2.2 |
| 1995 | 0.55 | 0.00 | 0.7 | 0.0 | 0.7 | 0.105 | 0.04 | 0.14 | 0.9 | 2.0 |
| 1996 | 3.00 | 0.00 | 4.1 | 0.0 | 4.1 | 0.106 | 0.01 | 0.11 | 4.6 | 2.6 |
| 1997 | 0.75 | 0.00 | 1.0 | 0.0 | 1.0 | 0.090 | 0.01 | 0.10 | 1.1 | 2.4 |
| 1998 | 1.48 | 0.00 | 2.0 | 0.0 | 2.0 | 0.105 | 0.01 | 0.11 | 2.2 | 2.1 |
| 1999 | 2.98 | 0.00 | 4.0 | 0.0 | 4.0 | 0.090 | 0.01 | 0.10 | 4.5 | 2.7 |
| 2000 | 2.80 | 0.00 | 3.8 | 0.0 | 3.8 | 0.079 | 0.01 | 0.09 | 4.1 | 2.9 |


| Basin: John Day |
| :---: |
| Population: Upper NF John Day |
| Sub-population: |
| Monitoring sites: Beaver Cr |
| Olive $\mathbf{C r}$ |
| Method: Redd Surveys |


|  | Critical Threshold |
| :--- | :--- |
| Viable Threshold | 0.12 |
|  | 0.46 |
|  |  |


| Females per Redd | $=0.81$ |
| ---: | :--- |
| females in spng pop. | $=0.60$ |



| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre-Harv $\begin{array}{r}6 \text { year } \\ \text {-Harv Wild Fi Moving Av }\end{array}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild fish | Hatch Fish | Wild Spwnrs | Hatch Spnrs |  | Columbia | John Day | Combined |  |  |
| 1977 | 3.0 | 0.0 | 4.1 | 0.0 | 4.1 | 0.078 | 0.12 | 0.19 | 5.0 |  |
| 1978 | 2.3 | 0.0 | 3.0 | 0.0 | 3.0 | 0.208 | 0.12 | 0.30 | 4.4 |  |
| 1979 | 1.5 | 0.0 | 2.0 | 0.0 | 2.0 | 0.196 | 0.12 | 0.29 | 2.9 |  |
| 1980 | 2.3 | 0.0 | 3.0 | 0.0 | 3.0 | 0.079 | 0.12 | 0.19 | 3.7 |  |
| 1981 | 2.0 | 0.0 | 2.7 | 0.0 | 2.7 | 0.087 | 0.12 | 0.20 | 3.4 |  |
| 1982 | 2.5 | 0.0 | 3.4 | 0.0 | 3.4 | 0.069 | 0.12 | 0.18 | 4.2 | 3.9 |
| 1983 | 3.8 | 0.0 | 5.1 | 0.0 | 5.1 | 0.069 | 0.12 | 0.18 | 6.2 | 4.1 |
| 1984 | 0.9 | 0.0 | 1.2 | 0.0 | 1.2 | 0.088 | 0.12 | 0.20 | 1.5 | 3.6 |
| 1985 | 4.8 | 0.0 | 6.4 | 0.0 | 6.4 | 0.121 | 0.12 | 0.23 | 8.3 | 4.5 |
| 1986 | 6.8 | 0.0 | 9.1 | 0.0 | 9.1 | 0.209 | 0.12 | 0.30 | 13.1 | 6.1 |
| 1987 | 4.0 | 0.0 | 5.4 | 0.0 | 5.4 | 0.139 | 0.12 | 0.24 | 7.1 | 6.7 |
| 1988 | 7.5 | 0.0 | 10.1 | 0.0 | 10.1 | 0.158 | 0.12 | 0.26 | 13.7 | 8.3 |
| 1989 | 2.3 | 0.0 | 3.0 | 0.0 | 3.0 | 0.172 | 0.12 | 0.27 | 4.2 | 8.0 |
| 1990 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.161 | 0.12 | 0.26 | 0.1 | 7.7 |
| 1991 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.160 | 0.04 | 0.19 | 0.1 | 6.4 |
| 1992 | 3.8 | 0.0 | 5.1 | 0.0 | 5.1 | 0.147 | 0.12 | 0.25 | 6.7 | 5.3 |
| 1993 | 2.0 | 0.0 | 2.7 | 0.0 | 2.7 | 0.164 | 0.12 | 0.26 | 3.7 | 4.7 |
| 1994 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.155 | 0.04 | 0.19 | 0.1 | 2.5 |
| 1995 | 1.3 | 0.0 | 1.7 | 0.0 | 1.7 | 0.105 | 0.04 | 0.14 | 2.0 | 2.1 |
| 1996 | 3.8 | 0.0 | 5.1 | 0.0 | 5.1 | 0.106 | 0.01 | 0.11 | 5.7 | 3.0 |
| 1997 | 1.4 | 0.0 | 1.8 | 0.0 | 1.8 | 0.090 | 0.01 | 0.10 | 2.0 | 3.4 |
| 1998 | 2.5 | 0.0 | 3.4 | 0.0 | 3.4 | 0.105 | 0.01 | 0.11 | 3.8 | 2.9 |
| 1999 | 1.6 | 0.0 | 2.1 | 0.0 | 2.1 | 0.090 | 0.01 | 0.10 | 2.4 | 2.7 |
| 2000 | 1.6 | 0.0 | 2.1 | 0.0 | 2.1 | 0.079 | 0.01 | 0.09 | 2.4 | 3.0 |


| Basin: John Day |
| :---: |
| Population: MF John Day |
| Sub-population: |
| Monitoring sites: Camp Cr <br> (primary) $\quad$ Lick Cr <br>  <br> Method: Redd Surveys <br>  |


|  |  |
| :--- | :--- |
| Critical Threshold | 0.83 |
| Viable Threshold | 2.24 |
|  | 4.80 |

Females per Redd
Prop. of females in spng pop.
$=0.81$

Average Distribution of Ages in return year

| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre-Harv | 6 -year |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Sprrs |  | Columbia | John Day | Combined | e-Haru Wild Fi | Moving Av |
| 1974 | 6.0 | 0.0 | 8.1 | 0.0 | 8.1 | 0.289 | 0.12 | 0.37 | 12.9 |  |
| 1975 | 9.5 | 0.0 | 12.8 | 0.0 | 12.8 | 0.128 | 0.12 | 0.23 | 16.7 |  |
| 1976 | 10.5 | 0.0 | 14.2 | 0.0 | 14.2 | 0.067 | 0.12 | 0.18 | 17.3 |  |
| 1977 | 16.8 | 0.0 | 22.7 | 0.0 | 22.7 | 0.078 | 0.12 | 0.19 | 27.9 |  |
| 1978 | 12.5 | 0.0 | 16.9 | 0.0 | 16.9 | 0.208 | 0.12 | 0.30 | 24.2 |  |
| 1979 | 2.0 | 0.0 | 2.7 | 0.0 | 2.7 | 0.196 | 0.12 | 0.29 | 3.9 | 17.2 |
| 1980 | 4.3 | 0.0 | 5.9 | 0.0 | 5.9 | 0.079 | 0.12 | 0.19 | 7.2 | 16.2 |
| 1981 | 7.3 | 0.0 | 9.9 | 0.0 | 9.9 | 0.087 | 0.12 | 0.20 | 12.3 | 15.5 |
| 1982 | 5.6 | 0.0 | 7.6 | 0.0 | 7.6 | 0.069 | 0.12 | 0.18 | 9.2 | 14.1 |
| 1983 | 3.2 | 0.0 | 4.3 | 0.0 | 4.3 | 0.069 | 0.12 | 0.18 | 5.2 | 10.3 |
| 1984 | 3.1 | 0.0 | 4.2 | 0.0 | 4.2 | 0.088 | 0.12 | 0.20 | 5.2 | 7.2 |
| 1985 | 11.0 | 0.0 | 14.9 | 0.0 | 14.9 | 0.121 | 0.12 | 0.23 | 19.2 | 9.7 |
| 1986 | 17.2 | 0.0 | 23.3 | 0.0 | 23.3 | 0.209 | 0.12 | 0.30 | 33.5 | 14.1 |
| 1987 | 13.2 | 0.0 | 17.8 | 0.0 | 17.8 | 0.139 | 0.12 | 0.24 | 23.5 | 16.0 |
| 1988 | 16.8 | 0.0 | 22.7 | 0.0 | 22.7 | 0.158 | 0.12 | 0.26 | 30.6 | 19.5 |
| 1989 | 8.6 | 0.0 | 11.6 | 0.0 | 11.6 | 0.172 | 0.12 | 0.27 | 15.9 | 21.3 |
| 1990 | 3.4 | 0.0 | 4.6 | 0.0 | 4.6 | 0.161 | 0.12 | 0.26 | 6.3 | 21.5 |
| 1991 | 8.4 | 0.0 | 11.3 | 0.0 | 11.3 | 0.160 | 0.04 | 0.19 | 14.1 | 20.6 |
| 1992 | 11.8 | 0.0 | 15.9 | 0.0 | 15.9 | 0.147 | 0.12 | 0.25 | 21.2 | 18.6 |
| 1993 | 3.8 | 0.0 | 5.1 | 0.0 | 5.1 | 0.164 | 0.12 | 0.26 | 6.9 | 15.8 |
| 1994 | 6.1 | 0.0 | 8.2 | 0.0 | 8.2 | 0.155 | 0.04 | 0.19 | 10.1 | 12.4 |
| 1995 | 2.0 | 0.0 | 2.6 | 0.0 | 2.6 | 0.105 | 0.04 | 0.14 | 3.1 | 10.3 |
| 1996 | 2.4 | 0.0 | 3.2 | 0.0 | 3.2 | 0.106 | 0.01 | 0.11 | 3.6 | 9.8 |
| 1997 | 2.2 | 0.0 | 3.0 | 0.0 | 3.0 | 0.090 | 0.01 | 0.10 | 3.3 | 8.0 |
| 1998 | 1.7 | 0.0 | 2.3 | 0.0 | 2.3 | 0.105 | 0.01 | 0.11 | 2.6 | 4.9 |
| 1999 | 5.2 | 0.0 | 7.0 | 0.0 | 7.0 | 0.090 | 0.01 | 0.10 | 7.7 | 5.1 |
| 2000 | 5.8 | 0.0 | 7.8 | 0.0 | 7.8 | 0.079 | 0.01 | 0.09 | 8.5 | 4.8 |


| Basin: John Day |
| :---: |
| Population: |
| SF John Day |
| Sub-population: <br> Monitoring sites: <br> (primary) <br> Uper Cr Murder's Cr <br> Tex Cr |
| Method: Redd Surveys |


|  | Critical Threshold |
| :--- | :--- |
| Viable Threshold | 0.55 |
|  | 1.67 |
|  | $\mathbf{2 . 6 3}$ |

$\begin{aligned} \text { Females per Redd } & =0.81 \\ \text { Prop. of females in spng pop. } & =0.60\end{aligned}$

| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre- Harv e-Harv Wild Fi | 6-year Moving Av |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Spnrs |  | Columbia | John Day | Combined |  |  |
| 1974 | 9.7 | 0.0 | 13.1 | 0.0 | 13.1 | 0.289 | 0.12 | 0.37 | 20.9 |  |
| 1975 | 13.9 | 0.0 | 18.8 | 0.0 | 18.8 | 0.128 | 0.12 | 0.23 | 24.5 |  |
| 1976 | 7.7 | 0.0 | 10.4 | 0.0 | 10.4 | 0.067 | 0.12 | 0.18 | 12.7 |  |
| 1977 | 9.4 | 0.0 | 12.7 | 0.0 | 12.7 | 0.078 | 0.12 | 0.19 | 15.6 |  |
| 1978 | 5.4 | 0.0 | 7.3 | 0.0 | 7.3 | 0.208 | 0.12 | 0.30 | 10.5 |  |
| 1979 | 2.8 | 0.0 | 3.8 | 0.0 | 3.8 | 0.196 | 0.12 | 0.29 | 5.3 | 14.9 |
| 1980 | 5.3 | 0.0 | 7.2 | 0.0 | 7.2 | 0.079 | 0.12 | 0.19 | 8.8 | 12.9 |
| 1981 | 4.2 | 0.0 | 5.7 | 0.0 | 5.7 | 0.087 | 0.12 | 0.20 | 7.1 | 10.0 |
| 1982 | 7.3 | 0.0 | 9.9 | 0.0 | 9.9 | 0.069 | 0.12 | 0.18 | 12.0 | 9.9 |
| 1983 | 8.9 | 0.0 | 12.0 | 0.0 | 12.0 | 0.069 | 0.12 | 0.18 | 14.7 | 9.7 |
| 1984 | 6.0 | 0.0 | 8.1 | 0.0 | 8.1 | 0.088 | 0.12 | 0.20 | 10.1 | 9.7 |
| 1985 | 11.4 | 0.0 | 15.4 | 0.0 | 15.4 | 0.121 | 0.12 | 0.23 | 19.9 | 12.1 |
| 1986 | 10.2 | 0.0 | 13.8 | 0.0 | 13.8 | 0.209 | 0.12 | 0.30 | 19.8 | 13.9 |
| 1987 | 13.6 | 0.0 | 18.4 | 0.0 | 18.4 | 0.139 | 0.12 | 0.24 | 24.2 | 16.8 |
| 1988 | 14.4 | 0.0 | 19.4 | 0.0 | 19.4 | 0.158 | 0.12 | 0.26 | 26.2 | 19.1 |
| 1989 | 2.6 | 0.0 | 3.5 | 0.0 | 3.5 | 0.172 | 0.12 | 0.27 | 4.8 | 17.5 |
| 1990 | 6.2 | 0.0 | 8.4 | 0.0 | 8.4 | 0.161 | 0.12 | 0.26 | 11.3 | 17.7 |
| 1991 | 3.1 | 0.0 | 4.2 | 0.0 | 4.2 | 0.160 | 0.04 | 0.19 | 5.2 | 15.3 |
| 1992 | 4.0 | 0.0 | 5.4 | 0.0 | 5.4 | 0.147 | 0.12 | 0.25 | 7.2 | 13.2 |
| 1993 | 2.4 | 0.0 | 3.2 | 0.0 | 3.2 | 0.164 | 0.12 | 0.26 | 4.4 | 9.9 |
| 1994 | 4.3 | 0.0 | 5.8 | 0.0 | 5.8 | 0.155 | 0.04 | 0.19 | 7.2 | 6.7 |
| 1995 | 2.1 | 0.0 | 2.8 | 0.0 | 2.8 | 0.105 | 0.04 | 0.14 | 3.3 | 6.4 |
| 1996 | 2.3 | 0.0 | 3.1 | 0.0 | 3.1 | 0.106 | 0.01 | 0.11 | 3.5 | 5.1 |
| 1997 | 1.4 | 0.0 | 1.9 | 0.0 | 1.9 | 0.090 | 0.01 | 0.10 | 2.1 | 4.6 |
| 1998 | 0.9 | 0.0 | 1.2 | 0.0 | 1.2 | 0.105 | 0.01 | 0.11 | 1.4 | 3.6 |
| 1999 | 0.9 | 0.0 | 1.2 | 0.0 | 1.2 | 0.090 | 0.01 | 0.10 | 1.3 | 3.1 |
| 2000 | 2.8 | 0.0 | 3.8 | 0.0 | 3.8 | 0.079 | 0.01 | 0.09 | 4.2 | 2.6 |



| Critical Threshold | 0.24 |
| :--- | :--- |
|  | Viable Threshold |
| Last 6-yr Average | 0.79 |
|  |  |

Females per Redd
Prop. of females in spng pop.

| $=0.81$ |
| :--- |
| $=0.60$ |


| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective <br> Spawners | Harvest Rates |  |  | Pre- Harv e-Harv Wild Fi | 6-year <br> Moving Av |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Spnrs |  | Columbia | John Day | Combined |  |  |
| 1974 | 4.0 | 0.0 | 5.3 | 0.0 | 5.3 | 0.289 | 0.12 | 0.37 | 8.5 |  |
| 1975 | 6.7 | 0.0 | 9.0 | 0.0 | 9.0 | 0.128 | 0.12 | 0.23 | 11.7 |  |
| 1976 | 3.6 | 0.0 | 4.9 | 0.0 | 4.9 | 0.067 | 0.12 | 0.18 | 6.0 |  |
| 1977 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.078 | 0.12 | 0.19 | 0.1 |  |
| 1978 | 4.3 | 0.0 | 5.9 | 0.0 | 5.9 | 0.208 | 0.12 | 0.30 | 8.4 |  |
| 1979 | 0.2 | 0.0 | 0.2 | 0.0 | 0.2 | 0.196 | 0.12 | 0.29 | 0.3 | 5.8 |
| 1980 | 4.3 | 0.0 | 5.7 | 0.0 | 5.7 | 0.079 | 0.12 | 0.19 | 7.1 | 5.6 |
| 1981 | 3.9 | 0.0 | 5.3 | 0.0 | 5.3 | 0.087 | 0.12 | 0.20 | 6.6 | 4.7 |
| 1982 | 1.8 | 0.0 | 2.4 | 0.0 | 2.4 | 0.069 | 0.12 | 0.18 | 2.9 | 4.2 |
| 1983 | 2.5 | 0.0 | 3.4 | 0.0 | 3.4 | 0.069 | 0.12 | 0.18 | 4.1 | 4.9 |
| 1984 | 3.5 | 0.0 | 4.7 | 0.0 | 4.7 | 0.088 | 0.12 | 0.20 | 5.9 | 4.5 |
| 1985 | 5.9 | 0.0 | 8.0 | 0.0 | 8.0 | 0.121 | 0.12 | 0.23 | 10.3 | 6.1 |
| 1986 | 11.1 | 0.0 | 15.0 | 0.0 | 15.0 | 0.209 | 0.12 | 0.30 | 21.5 | 8.5 |
| 1987 | 20.7 | 0.0 | 27.9 | 0.0 | 27.9 | 0.139 | 0.12 | 0.24 | 36.8 | 13.6 |
| 1988 | 16.3 | 0.0 | 22.1 | 0.0 | 22.1 | 0.158 | 0.12 | 0.26 | 29.8 | 18.1 |
| 1989 | 3.3 | 0.0 | 4.5 | 0.0 | 4.5 | 0.172 | 0.12 | 0.27 | 6.2 | 18.4 |
| 1990 | 3.5 | 0.0 | 4.7 | 0.0 | 4.7 | 0.161 | 0.12 | 0.26 | 6.4 | 18.5 |
| 1991 | 3.4 | 0.0 | 4.6 | 0.0 | 4.6 | 0.160 | 0.04 | 0.19 | 5.7 | 17.7 |
| 1992 | 2.3 | 0.0 | 3.2 | 0.0 | 3.2 | 0.147 | 0.12 | 0.25 | 4.2 | 14.8 |
| 1993 | 2.0 | 0.0 | 2.7 | 0.0 | 2.7 | 0.164 | 0.12 | 0.26 | 3.7 | 9.3 |
| 1994 | 0.4 | 0.0 | 0.5 | 0.0 | 0.5 | 0.155 | 0.04 | 0.19 | 0.6 | 4.5 |
| 1995 | 0.5 | 0.0 | 0.7 | 0.0 | 0.7 | 0.105 | 0.04 | 0.14 | 0.8 | 3.6 |
| 1996 | 2.1 | 0.0 | 2.8 | 0.0 | 2.8 | 0.106 | 0.01 | 0.11 | 3.1 | 3.0 |
| 1997 | 1.35 | 0.0 | 1.8 | 0.0 | 1.8 | 0.090 | 0.01 | 0.10 | 2.0 | 2.4 |
| 1998 | 1.45 | 0.0 | 2.0 | 0.0 | 2.0 | 0.105 | 0.01 | 0.11 | 2.2 | 2.1 |
| 1999 | 2.35 | 0.0 | 3.2 | 0.0 | 3.2 | 0.090 | 0.01 | 0.10 | 3.5 | 2.0 |
| 2000 | 3.0 | 0.0 | 4.1 | 0.0 | 4.1 | 0.079 | 0.01 | 0.09 | 4.4 | 2.7 |


| Basin: John Day |
| :---: |
| Population: Upper Mainstem JD |
| Sub-population: |
| Monitoring sites: Canyon Cr |
| (primary) |
| Fields $\mathbf{C r}$ |
| Riley Cr |
| Bear (Grant Co.) |
| Beech $\mathbf{~ C r}$ |
| EF Beech Cr |
| McClellan Cr |
| Method: |
| Redd Surveys |



Average Distribution of Ages in return year

| Females per Redd | $=0.81$ |
| ---: | :--- |
| Prop. of females in spng pop. | $=0.60$ |


| Spawning Year | Redds/Mile |  | Fish/Mile |  | Effective Spawners | Harvest Rates |  |  | Pre- Harv 6 - yeare-Harv Wild Fi Moving Av |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild Fish | Hatch Fish | Wild Spwnrs | Hatch Spnrs |  | Columbia | John Day | Combined |  |  |
| 1974 | 5.1 | 0.0 | 6.9 | 0.0 | 6.9 | 0.289 | 0.12 | 0.37 | 10.9 |  |
| 1975 | 6.8 | 0.0 | 9.2 | 0.0 | 9.2 | 0.128 | 0.12 | 0.23 | 12.0 |  |
| 1976 | 5.5 | 0.0 | 7.4 | 0.0 | 7.4 | 0.067 | 0.12 | 0.18 | 9.1 |  |
| 1977 | 8.4 | 0.0 | 11.3 | 0.0 | 11.3 | 0.078 | 0.12 | 0.19 | 14.0 |  |
| 1978 | 4.6 | 0.0 | 6.2 | 0.0 | 6.2 | 0.208 | 0.12 | 0.30 | 8.8 |  |
| 1979 | 0.9 | 0.0 | 1.3 | 0.0 | 1.3 | 0.196 | 0.12 | 0.29 | 1.8 | 9.4 |
| 1980 | 5.5 | 0.0 | 7.4 | 0.0 | 7.4 | 0.079 | 0.12 | 0.19 | 9.1 | 9.1 |
| 1981 | 3.2 | 0.0 | 4.4 | 0.0 | 4.4 | 0.087 | 0.12 | 0.20 | 5.4 | 8.0 |
| 1982 | 3.9 | 0.0 | 5.3 | 0.0 | 5.3 | 0.069 | 0.12 | 0.18 | 6.5 | 7.6 |
| 1983 | 5.1 | 0.0 | 6.9 | 0.0 | 6.9 | 0.069 | 0.12 | 0.18 | 8.4 | 6.7 |
| 1984 | 3.3 | 0.0 | 4.5 | 0.0 | 4.5 | 0.088 | 0.12 | 0.20 | 5.6 | 6.1 |
| 1985 | 7.8 | 0.0 | 10.6 | 0.0 | 10.6 | 0.121 | 0.12 | 0.23 | 13.7 | 8.1 |
| 1986 | 16.2 | 0.0 | 21.9 | 0.0 | 21.9 | 0.209 | 0.12 | 0.30 | 31.5 | 11.8 |
| 1987 | 14.5 | 0.0 | 19.6 | 0.0 | 19.6 | 0.139 | 0.12 | 0.24 | 25.9 | 15.3 |
| 1988 | 15.7 | 0.0 | 21.2 | 0.0 | 21.2 | 0.158 | 0.12 | 0.26 | 28.6 | 18.9 |
| 1989 | 4.8 | 0.0 | 6.4 | 0.0 | 6.4 | 0.172 | 0.12 | 0.27 | 8.8 | 19.0 |
| 1990 | 5.4 | 0.0 | 7.2 | 0.0 | 7.2 | 0.161 | 0.12 | 0.26 | 9.8 | 19.7 |
| 1991 | 3.4 | 0.0 | 4.6 | 0.0 | 4.6 | 0.160 | 0.04 | 0.19 | 5.7 | 18.4 |
| 1992 | 7.7 | 0.0 | 10.3 | 0.0 | 10.3 | 0.147 | 0.12 | 0.25 | 13.8 | 15.4 |
| 1993 | 2.9 | 0.0 | 3.9 | 0.0 | 3.9 | 0.164 | 0.12 | 0.26 | 5.3 | 12.0 |
| 1994 | 4.2 | 0.0 | 5.7 | 0.0 | 5.7 | 0.155 | 0.04 | 0.19 | 7.1 | 8.4 |
| 1995 | 1.1 | 0.0 | 1.4 | 0.0 | 1.4 | 0.105 | 0.04 | 0.14 | 1.7 | 7.2 |
| 1996 | 1.6 | 0.0 | 2.2 | 0.0 | 2.2 | 0.106 | 0.01 | 0.11 | 2.5 | 6.0 |
| 1997 | 1.6 | 0.0 | 2.1 | 0.0 | 2.1 | 0.090 | 0.01 | 0.10 | 2.4 | 5.5 |
| 1998 | 2.7 | 0.0 | 3.6 | 0.0 | 3.6 | 0.105 | 0.01 | 0.11 | 4.1 | 3.8 |
| 1999 | 0.9 | 0.0 | 1.3 | 0.0 | 1.3 | 0.090 | 0.01 | 0.10 | 1.4 | 3.2 |
| 2000 | 2.5 | 0.0 | 3.4 | 0.0 | 3.4 | 0.079 | 0.01 | 0.09 | 3.7 | 2.6 |


| Basin: Deschutes |
| :---: |
| Population: Deschutes |
| Sub-population: |
| Monitoring sites: Sherars Falls |
| Methods: <br> Mark-recapture population estimate from <br> sub-sample of run tagged at Sherars Falls. <br> Recoveries at Pelton and Warm Springs <br> NFH Traps. |


| Critical Threshold | 473 |
| :--- | :--- |
| Viable Threshold | 1479 |
|  | 7395 |


| Spawning Year | Wild Escapm |  |  |  | Total | Pre- Han | Wild Abund <br> 6 -yr <br> Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Pop.Estm | Wild Spwnrs | RBH | Hatch Strays | Spawners | Wild Abund |  |
| 1978 | 6600 | 5808 | 3248 | 237 | 9293 | 7446 |  |
| 1979 | 2800 | 2464 | 1084 | 0 | 3548 | 3159 |  |
| 1980 | 4200 | 4150 | 2140 | 157 | 6447 | 4243 |  |
| 1981 | 4100 | 4051 | 2645 | 129 | 6825 | 4142 |  |
| 1982 | 6900 | 6817 | 1584 | 266 | 8668 | 6971 |  |
| 1983 | 6567 | 6488 | 1554 | 396 | 8439 | 6634 | 5432 |
| 1984 | 8228 | 8129 | 3941 | 1253 | 13323 | 8312 | 5577 |
| 1985 | 7721 | 7628 | 3377 | 551 | 11557 | 7800 | 6350 |
| 1986 | 9624 | 9509 | 3343 | 837 | 13688 | 9722 | 7264 |
| 1987 | 6207 | 6133 | 5336 | 1913 | 13381 | 6270 | 7618 |
| 1988 | 5367 | 5303 | 6620 | 2149 | 14072 | 5422 | 7360 |
| 1989 | 3546 | 3503 | 2140 | 724 | 6368 | 3582 | 6852 |
| 1990 | 4278 | 4227 | 1598 | 763 | 6588 | 4322 | 6186 |
| 1991 | 3653 | 3609 | 1145 | 604 | 5358 | 3690 | 5502 |
| 1992 | 4826 | 4768 | 1960 | 2111 | 8839 | 4875 | 4694 |
| 1993 | 904 | 893 | 1077 | 1073 | 3043 | 913 | 3801 |
| 1994 | 1487 | 1469 | 830 | 1228 | 3527 | 1502 | 3148 |
| 1995 | 482 | 476 | 814 | 904 | 2194 | 487 | 2632 |
| 1996 | 1662 | 1642 | 1383 | 3197 | 6222 | 1679 | 2191 |
| 1997 | 3458 | 3417 | 1615 | 6313 | 11344 | 3493 | 2158 |
| 1998 | 1820 | 1798 | 1997 | 3772 | 7568 | 1839 | 1652 |
| 1999 | 3800 | 3754 | 1969 | 2627 | 8350 | 3839 | 2140 |
| 2000 | 4790 | 4733 | 1263 | 4084 | 10079 | 4839 | 2696 |


| Basin: Deschutes |
| :--- |
| Population: Deschutes |
| Sub-population: Warm Springs |
| Monitoring sites: Warm Springs |
| NFH Trap |
| Method: <br> Total count of returning fish. No hatchery <br> fish allowed to pass into Warm Springs <br> River above the trap. |



Average Distribution of Ages in return year

| Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.05 | 0.00 | 0.15 | 0.41 | 0.33 | 0.07 |


| Spawning Year | Spawners |  | TotalSpawners | Hr above Shearars | Hr below Sherars | Columbia Mainst HR | overall <br> Harv Rate | Pre- Harv Wild Fish |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Wild | Hatchery |  |  |  |  |  |  |  |
| 1980 | 378 | 0.0 | 378 | 0.01 | 0.01 | 0.079 | 0.097 | 419 |  |
| 1981 | 311 | 0.0 | 311 | 0.01 | 0.01 | 0.087 | 0.105 | 348 |  |
| 1982 | 397 | 0.0 | 397 | 0.01 | 0.01 | 0.069 | 0.088 | 435 |  |
| 1983 | 569 | 0.0 | 569 | 0.01 | 0.01 | 0.069 | 0.088 | 624 |  |
| 1984 | 255 | 0.0 | 255 | 0.01 | 0.01 | 0.088 | 0.106 | 285 |  |
| 1985 | 431 | 0.0 | 431 | 0.01 | 0.01 | 0.121 | 0.138 | 500 | 435 |
| 1986 | 577 | 0.0 | 577 | 0.01 | 0.01 | 0.209 | 0.225 | 744 | 489 |
| 1987 | 373 | 0.0 | 373 | 0.01 | 0.01 | 0.139 | 0.156 | 442 | 505 |
| 1988 | 822 | 0.0 | 822 | 0.01 | 0.01 | 0.158 | 0.175 | 996 | 599 |
| 1989 | 522 | 0.0 | 522 | 0.01 | 0.01 | 0.172 | 0.188 | 643 | 602 |
| 1990 | 385 | 0.0 | 385 | 0.01 | 0.01 | 0.161 | 0.178 | 468 | 632 |
| 1991 | 339 | 0.0 | 339 | 0.01 | 0.01 | 0.160 | 0.177 | 412 | 618 |
| 1992 | 165 | 0.0 | 165 | 0.01 | 0.01 | 0.147 | 0.164 | 197 | 526 |
| 1993 | 280 | 0.0 | 280 | 0.01 | 0.01 | 0.164 | 0.181 | 342 | 510 |
| 1994 | 79 | 0.0 | 79 | 0.01 | 0.01 | 0.155 | 0.172 | 95 | 360 |
| 1995 | 135 | 0.0 | 135 | 0.01 | 0.01 | 0.105 | 0.123 | 154 | 278 |
| 1996 | 95 | 0.0 | 95 | 0.01 | 0.01 | 0.106 | 0.124 | 108 | 218 |
| 1997 | 85 | 0.0 | 85 | 0.01 | 0.01 | 0.090 | 0.108 | 95 | 165 |
| 1998 | 243 | 0.0 | 243 | 0.01 | 0.01 | 0.105 | 0.123 | 277 | 179 |
| 1999 | 214 | 0.0 | 214 | 0.01 | 0.01 | 0.090 | 0.108 | 240 | 162 |



| Basin: Hood River |
| :---: |
| Population: Hood River StS |
| Sub-population: |
| Monitoring sites: Powerdale Dam |
| Trap |
| Method: Total count of |
| returning fish. |




| Basin: Hood River |
| :---: |
| Population: Hood River Stw |
| Sub-population: |
| Monitoring sites: Powerdale Dam |
| Trap |
| Method: Total count of |
| returning fish. |


| Critical Threshold |
| :--- | :--- |
| Viable Threshold |
| Last 6-yr Average |



Average Distribution of Ages at time of spawning

| Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.05 | 0.00 | 0.07 | 0.60 | 0.25 | 0.02 |


| Spawning Year | Wild Fish |  | Hatchery Fish |  | Effective <br> Tot. Spwnrs | Wild Fish Harvest Rates |  |  | Pre- Harv <br> e-Harv Wild F | 6-yr <br> Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | counted | escapem | counted | escapem |  | Out-basin | In-basin | Combined |  |  |
| 1992 | 699 | 621 | 318 | 281 | 902 | 0.147 | 0.010 | 0.22 | 892 |  |
| 1993 | 412 | 343 | 238 | 11 | 354 | 0.164 | 0.010 | 0.22 | 527 |  |
| 1994 | 405 | 301 | 176 | 5 | 306 | 0.155 | 0.010 | 0.17 | 488 |  |
| 1995 | 206 | 161 | 111 | 5 | 166 | 0.105 | 0.010 | 0.12 | 235 |  |
| 1996 | 280 | 211 | 280 | 162 | 373 | 0.106 | 0.010 | 0.15 | 328 |  |
| 1997 | 289 | 239 | 641 | 254 | 493 | 0.090 | 0.010 | 0.14 | 336 | 468 |
| 1998 | 227 | 182 | 393 | 164 | 346 | 0.105 | 0.010 | 0.14 | 263 | 363 |
| 1999 | 301 | 258 | 323 | 200 | 458 | 0.090 | 0.010 | 0.14 | 349 | 333 |
| 2000 | 930 | 876 | 299 | 200 | 1076 | 0.079 | 0.010 | 0.14 | 1077 | 431 |


| Basin: Sandy |  |  |  | --Wild Abund - 0 - Moving Avg |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Population: Sandy |  |  |  |  |  |  |  |  |  |  |
| Sub-population: |  |  |  |  |  |  |  |  |  |  |
| Monitoring sites: Marmot Dam |  |  |  |  |  |  |  |  |  |  |
| Method: Counts estimated from video pictures at ladder until 1998. from 1998 to present fish have been trapped and directly counted. Likewise for identification hatchery fish; prior to 1998 identification was based upon run timing (hatchery fish earlier than wild). |  |  |  |  |  |  |  |  | $1000$ |  |
| Critical Threshold |  | 76 |  | Spawner Year |  |  |  |  |  |  |
| Viable Threshold |  | 336 |  |  |  |  |  |  |  |  |
| Last 6 | yr Average | 651 |  |  |  |  |  |  |  |  |
| Note: Wild fish removed for hatchery broodstock development. Also hatchery fish prevented from passing upstream of trap site starting in 1999. |  |  |  | Average Distribution of Ages in return year |  |  |  |  |  |  |
|  |  |  |  |  | Repeat |  |  |  | Age 5 | Age 6 |
|  |  |  |  |  | 0.11 | 0.00 | 0.01 | 0.63 | 0.23 | 0.02 |
| Spawning |  | wild fish |  | Effective |  | Wild Fish Harvest Rates |  |  | Pre-Han | 6-yr |
| Year | wild count | removed | SpwnrsWild | SpwnrsHatc | Tot. Spwnrs | Out-basin | In-basin | Combined | Wild Abund | Moving Avg |
| 1978 | 1342 |  | 1342 | 2729 | 4071 | 0.00 | 0.40 | 0.40 | 2237 |  |
| 1979 | 1344 |  | 1344 | 656 | 2000 | 0.00 | 0.40 | 0.40 | 2240 |  |
| 1980 | 2157 |  | 2157 | 858 | 3015 | 0.00 | 0.40 | 0.40 | 3595 |  |
| 1981 | 1338 |  | 1338 | 2740 | 4078 | 0.00 | 0.40 | 0.40 | 2230 |  |
| 1982 | 1602 |  | 1602 | 1087 | 2689 | 0.00 | 0.40 | 0.40 | 2670 |  |
| 1983 | 856 |  | 856 | 1593 | 2449 | 0.00 | 0.40 | 0.40 | 1427 | 2400 |
| 1984 | 1176 |  | 1176 | 1056 | 2232 | 0.00 | 0.40 | 0.40 | 1960 | 2354 |
| 1985 | 1505 |  | 1505 | 1336 | 2841 | 0.00 | 0.40 | 0.40 | 2508 | 2398 |
| 1986 | 1995 |  | 1995 | 757 | 2752 | 0.00 | 0.40 | 0.40 | 3325 | 2353 |
| 1987 | 1785 |  | 1785 | 1890 | 3675 | 0.00 | 0.40 | 0.40 | 2975 | 2478 |
| 1988 | 1401 |  | 1401 | 2039 | 3440 | 0.00 | 0.40 | 0.40 | 2335 | 2422 |
| 1989 | 1356 |  | 1356 | 1637 | 2993 | 0.00 | 0.40 | 0.40 | 2260 | 2561 |
| 1990 | 1438 |  | 1438 | 1627 | 3065 | 0.00 | 0.40 | 0.40 | 2397 | 2633 |
| 1991 | 707 |  | 707 | 1288 | 1995 | 0.00 | 0.40 | 0.40 | 1178 | 2412 |
| 1992 | 956 |  | 956 | 1962 | 2918 | 0.00 | 0.04 | 0.04 | 996 | 2023 |
| 1993 | 1008 |  | 1008 | 628 | 1636 | 0.00 | 0.04 | 0.04 | 1050 | 1703 |
| 1994 | 802 |  | 802 | 765 | 1567 | 0.00 | 0.04 | 0.04 | 835 | 1453 |
| 1995 | 653 |  | 653 | 1027 | 1680 | 0.00 | 0.04 | 0.04 | 680 | 1189 |
| 1996 | 220 |  | 220 | 316 | 536 | 0.00 | 0.04 | 0.04 | 229 | 828 |
| 1997 | 924 |  | 924 | 474 | 1398 | 0.00 | 0.04 | 0.040 | 963 | 792 |
| 1998 | 584 |  | 584 | 359 | 943 | 0.00 | 0.04 | 0.040 | 608 | 728 |
| 1999 | 629 |  | 629 | 0 | 629 | 0.00 | 0.04 | 0.040 | 655 | 662 |
| 2000 | 741 | 123 | 618 | 0 | 618 | 0.00 | 0.04 | 0.040 | 772 | 651 |



| Basin: Molalla |
| :---: |
| Population: Molalla |
| Sub-population: |
| Monitoring sites: Index sites |
|  |
| Method: Redd Surveys |



|  |  |  | Average Distribution of Ages at time of spawning |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
|  |  |  | 0.10 | 0.00 | 0.00 | 0.83 | 0.07 | 0.00 |
|  |  |  |  |  |  | Wild Abund |  |  |
| Spawning |  |  | Effective | Wild Fish | Pre- Harv | 6-yr |  |  |
| Year | SpwnrsWild | SpwnrsHatc | Tot. Spwnrs | Harv Rate | e-Harv Wild F | Moving Avg |  |  |
| 1980 | 41.1 | 35.0 | 76.1 | 0.21 | 52.0 |  |  |  |
| 1981 | 33.6 | 28.6 | 62.2 | 0.21 | 42.5 |  |  |  |
| 1982 | 29.5 | 25.1 | 54.6 | 0.21 | 37.3 |  |  |  |
| 1983 | 20.2 | 17.2 | 37.4 | 0.21 | 25.6 |  |  |  |
| 1984 | 28.5 | 24.3 | 52.8 | 0.21 | 36.1 |  |  |  |
| 1985 | 39.8 | 33.9 | 73.7 | 0.21 | 50.3 |  |  |  |
| 1986 | 34.5 | 29.4 | 63.9 | 0.21 | 43.7 | 39.3 |  |  |
| 1987 | 28.8 | 24.5 | 53.3 | 0.21 | 36.4 | 38.2 |  |  |
| 1988 | 35.0 | 29.9 | 64.9 | 0.21 | 44.4 | 39.4 |  |  |
| 1989 | 28.0 | 23.9 | 51.9 | 0.21 | 35.5 | 41.1 |  |  |
| 1990 | 23.8 | 20.3 | 44.1 | 0.21 | 30.1 | 40.1 |  |  |
| 1991 | 17.8 | 15.1 | 32.9 | 0.21 | 22.5 | 35.4 |  |  |
| 1992 | 24.0 | 7.2 | 31.1 | 0.02 | 24.5 | 32.2 |  |  |
| 1993 | 7.2 | 2.3 | 9.4 | 0.02 | 7.3 | 27.4 |  |  |
| 1994 | 29.3 | 9.3 | 38.6 | 0.02 | 29.9 | 25.0 |  |  |
| 1995 | 11.2 | 3.6 | 14.8 | 0.02 | 11.5 | 21.0 |  |  |
| 1996 | 5.1 | 1.6 | 6.6 | 0.02 | 5.2 | 16.8 |  |  |
| 1997 | 7.5 | 2.4 | 9.9 | 0.02 | 7.7 | 14.3 |  |  |
| 1998 | 17.9 | 5.4 | 23.3 | 0.02 | 18.3 | 13.3 |  |  |
| 1999 | 13.9 | 4.1 | 18.0 | 0.02 | 14.1 | 14.4 |  |  |
| 2000 | 24.1 | 0.5 | 24.6 | 0.02 | 24.6 | 13.6 |  |  |


| Basin: Santiam |
| :---: |
| Population: North Santiam |
| Sub-population: |
| Monitoring sites: Index Sites |
|  |
| Method: Redd counts |


| Critical Threshold | 9.8 |
| :--- | ---: |
|  | Viable Threshold |
|  | 16.4 |
|  | 21.2 |


Average Distribution of Ages at time of spawning

| Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.10 | 0.00 | 0.00 | 0.83 | 0.07 | 0.00 |


| Spawning Year | SpwnrsWild | SpwnrsHatc | Effective <br> Tot. Spwnrs | Wild Fish Harv Rate | Pre- Harv e-harv Wild Fi | 6-yT <br> Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1983 | 43.1 | 7.5 | 50.5 | 0.21 | 54.5 |  |
| 1984 | 46.7 | 8.1 | 54.8 | 0.21 | 59.1 |  |
| 1985 | 71.7 | 12.4 | 84.1 | 0.21 | 90.7 |  |
| 1986 | 49.6 | 8.6 | 58.2 | 0.21 | 62.8 |  |
| 1987 | 46.5 | 8.1 | 54.6 | 0.21 | 58.9 |  |
| 1988 | 25.1 | 4.4 | 29.5 | 0.21 | 31.8 | 59.6 |
| 1989 | 48.6 | 8.4 | 57.1 | 0.21 | 61.5 | 60.8 |
| 1990 | 44.1 | 7.7 | 51.8 | 0.21 | 55.9 | 60.3 |
| 1991 | 33.3 | 5.8 | 39.1 | 0.21 | 42.2 | 52.2 |
| 1992 | 25.3 | 4.4 | 29.7 | 0.02 | 25.8 | 46.0 |
| 1993 | 22.4 | 4.4 | 26.7 | 0.02 | 22.8 | 40.0 |
| 1994 | 27.2 | 4.1 | 31.3 | 0.02 | 27.7 | 39.3 |
| 1995 | 6.7 | 0.8 | 7.5 | 0.02 | 6.8 | 30.2 |
| 1996 | 11.8 | 1.5 | 13.3 | 0.02 | 12.0 | 22.9 |
| 1997 | 23.2 | 2.3 | 25.4 | 0.02 | 23.6 | 19.8 |
| 1998 | 24.2 | 10.1 | 34.2 | 0.02 | 24.7 | 19.6 |
| 1999 | 31.0 | 11.2 | 42.2 | 0.02 | 31.6 | 21.1 |
| 2000 | 27.9 | 3.9 | 31.8 | 0.02 | 28.4 | 21.2 |


| Basin: Santiam |
| :---: |
| Population: Upper S. Santiam |
| Sub-population: |
| Monitoring sites: Foster Dam <br> Trap |
| Method: Total count of |
| returning fish. |


|  | Critical Threshold |
| :--- | ---: |
| Viable Threshold | 30 |
|  | 108 |
|  |  |



Average Distribution of Ages at time of spawning

| Repeat Age 2  Age 3 Age 4  <br> 0.10 0.00 0.00 0.83 0.07 0.00 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |


| Spawning Year | SpwnrsWild | SpwnrsHatc | Effective Tot. Spwnrs | Wild Fish Harv Rate | $\begin{gathered} \text { Pre-Harv } \\ \text { e-harv Wild Fi } \end{gathered}$ | $6-y r$ Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1973 | 755 | 0 | 755 | 0.21 | 956 |  |
| 1974 | 695 | 0 | 695 | 0.21 | 880 |  |
| 1975 | 354 | 0 | 354 | 0.21 | 448 |  |
| 1976 | 302 | 0 | 302 | 0.21 | 382 |  |
| 1977 | 503 | 0 | 503 | 0.21 | 637 |  |
| 1978 | 488 | 0 | 488 | 0.21 | 618 | 653 |
| 1979 | 149 | 0 | 149 | 0.21 | 189 | 526 |
| 1980 | 515 | 0 | 515 | 0.21 | 652 | 488 |
| 1981 | 317 | 0 | 317 | 0.21 | 401 | 480 |
| 1982 | 234 | 165 | 399 | 0.21 | 296 | 465 |
| 1983 | 134 | 66 | 200 | 0.21 | 170 | 388 |
| 1984 | 504 | 993 | 1497 | 0.21 | 638 | 391 |
| 1985 | 355 | 629 | 984 | 0.21 | 449 | 434 |
| 1986 | 326 | 485 | 811 | 0.21 | 413 | 395 |
| 1987 | 214 | 253 | 467 | 0.21 | 271 | 373 |
| 1988 | 656 | 423 | 1079 | 0.18 | 800 | 457 |
| 1989 | 222 | 62 | 284 | 0.18 | 271 | 474 |
| 1990 | 272 | 10 | 282 | 0.18 | 332 | 423 |
| 1991 | 139 | 0 | 139 | 0.18 | 170 | 376 |
| 1992 | 361 | 0 | 361 | 0.03 | 372 | 369 |
| 1993 | 256 | 0 | 256 | 0.03 | 264 | 368 |
| 1994 | 234 | 0 | 234 | 0.02 | 239 | 274 |
| 1995 | 297 | 0 | 297 | 0.02 | 303 | 280 |
| 1996 | 131 | 0 | 131 | 0.02 | 134 | 247 |
| 1997 | 336 | 0 | 336 | 0.02 | 343 | 276 |
| 1998 | 359 | 0 | 359 | 0.02 | 366 | 275 |
| 1999 | 328 | 0 | 328 | 0.02 | 335 | 287 |
| 2000 | 326 | 0 | 326 | 0.02 | 333 | 302 |


| Basin: Santiam |
| :---: |
| Population: Lower S. Santiam |
| Sub-population: |
| Monitoring sites: Index sites |
| Method: Redd Counts |


| Critical Threshold | 2.10 |
| :---: | :---: |
| Viable Threshold | 8.10 |
| Last 6-yr Average | 8.09 |

Average Distribution of Ages at time of spawning

| Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.10 | 0.00 | 0.00 | 0.83 | 0.07 | 0.00 |


| Spawning Year | SpwnrsWild | SpwnrsHatc | Effective Tot. Spwnirs | Wild Fish Harv Rate | Pre- Harv e-harv Wild Fi | $6-y r$ Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1983 | 16.8 | 8.3 | 25 | 0.21 | 21.3 |  |
| 1984 | 11.1 | 21.9 | 33 | 0.21 | 14.1 |  |
| 1985 | 17.2 | 30.4 | 48 | 0.21 | 21.7 |  |
| 1986 | 14.2 | 21.2 | 35 | 0.21 | 18.0 |  |
| 1987 | 15.5 | 18.3 | 34 | 0.21 | 19.6 |  |
| 1988 | 19.8 | 12.8 | 33 | 0.21 | 25.1 |  |
| 1989 | 17.1 | 4.8 | 22 | 0.21 | 21.6 | 20.0 |
| 1990 | 30.0 | 1.1 | 31 | 0.21 | 38.0 | 24.0 |
| 1991 | 33.7 | 0.0 | 34 | 0.21 | 42.7 | 27.5 |
| 1992 | 29.5 | 0.0 | 30 | 0.02 | 30.1 | 29.5 |
| 1993 | 16.0 | 0.0 | 16 | 0.02 | 16.3 | 29.0 |
| 1994 | 25.6 | 0.0 | 26 | 0.02 | 26.1 | 29.1 |
| 1995 | 8.5 | 0.0 | 8 | 0.02 | 8.6 | 27.0 |
| 1996 | 3.9 | 0.0 | 4 | 0.02 | 4.0 | 21.3 |
| 1997 | 9.9 | 0.0 | 10 | 0.02 | 10.1 | 15.9 |
| 1998 | 10.6 | 0.0 | 11 | 0.02 | 10.8 | 12.7 |
| 1999 | 4.1 | 0.0 | 4 | 0.02 | 4.2 | 10.6 |
| 2000 | 10.6 | 0.0 | 11 | 0.02 | 10.8 | 8.1 |


| Basin: Calapooia |
| :---: |
| Population: Calapooia |
| Sub-population: |
| Monitoring sites: Index sites |
|  |
| Method: Redd Surveys |



| Critical Threshold | 0.70 |
| :---: | :---: |
| Viable Threshold | 2.20 |
|  | $\mathbf{B . 2 5}$ |

Average Distribution of Ages at time of spawning


| Basin: Nehalem |
| :---: |
| Population: Salmonberry |
| Sub-population: |
| Monitoring sites: Enright Mainstem |
| Site |
|  |
| Method: Spawning Survey <br> Counts of Observed Fish |


|  |  |
| :--- | :--- |
| Critical Threshold | 0.44 |
| Viable Threshold | 1.44 |
|  | $\mathbf{4 . 8 0}$ |


Average Distribution of Ages at time of spawning

| Repeat | Age 2 | Age 3 | Age 4 |  | Age 5 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.15 | 0.00 | 0.09 | 0.64 | 0.11 | 0.00 |


| Spawning Year | SpwnrsWild | SpwrrsHatc | Effective Tot. Spwnrs | Wild Fish Harv Rate | Pre- Han <br> Abundance | $6-y r$ Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1973 | 12.3 | 0.0 | 12.3 | 0.15 | 14.5 |  |
| 1974 | 5.7 | 0.0 | 5.7 | 0.15 | 6.7 |  |
| 1975 | 31.0 | 0.0 | 31.0 | 0.15 | 36.5 |  |
| 1976 | 10.7 | 0.0 | 10.7 | 0.15 | 12.5 |  |
| 1977 | 9.7 | 0.0 | 9.7 | 0.15 | 11.4 |  |
| 1978 | 11.6 | 0.0 | 11.6 | 0.15 | 13.7 | 15.9 |
| 1979 | 9.3 | 0.0 | 9.3 | 0.15 | 11.0 | 15.3 |
| 1980 | 13.7 | 0.0 | 13.7 | 0.15 | 16.1 | 16.9 |
| 1981 | 8.0 | 0.0 | 8.0 | 0.15 | 9.4 | 12.3 |
| 1982 | 5.0 | 0.0 | 5.0 | 0.15 | 5.9 | 11.2 |
| 1983 | 2.3 | 0.0 | 2.3 | 0.15 | 2.7 | 9.8 |
| 1984 | 4.0 | 0.0 | 4.0 | 0.15 | 4.7 | 8.3 |
| 1985 | 15.0 | 0.0 | 15.0 | 0.15 | 17.6 | 9.4 |
| 1986 | 7.7 | 0.0 | 7.7 | 0.15 | 9.0 | 8.2 |
| 1987 | 15.3 | 0.0 | 15.3 | 0.15 | 18.0 | 9.7 |
| 1988 | 9.3 | 0.0 | 9.3 | 0.15 | 11.0 | 10.5 |
| 1989 | 6.7 | 0.0 | 6.7 | 0.15 | 7.8 | 11.4 |
| 1990 | 14.7 | 0.0 | 14.7 | 0.15 | 17.3 | 13.5 |
| 1991 | 5.0 | 0.0 | 5.0 | 0.15 | 5.9 | 11.5 |
| 1992 | 6.0 | 0.0 | 6.0 | 0.02 | 6.1 | 11.0 |
| 1993 | 6.3 | 0.0 | 6.3 | 0.02 | 6.5 | 9.1 |
| 1994 | 6.0 | 0.0 | 6.0 | 0.02 | 6.1 | 8.3 |
| 1995 | 9.0 | 0.0 | 9.0 | 0.02 | 9.2 | 8.5 |
| 1997 | 5.7 | 0.0 | 5.7 | 0.02 | 5.8 | 6.6 |
| 1997 | 3.7 | 0.0 | 3.7 | 0.02 | 3.7 | 6.2 |
| 1998 | 1.7 | 0.0 | 1.7 | 0.02 | 1.7 | 5.5 |
| 1999 | 2.3 | 0.0 | 2.3 | 0.02 | 2.3 | 4.8 |




|  |  |  |  | Average D | ribution | Ages at ti | of spaw |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |  |
|  |  |  |  | 0.14 | 0.00 | 0.00 | 0.59 | 0.25 | 0.03 |  |
| SpwnrsWild | SpwnrsHatc | Effective <br> Tot. Spwnrs | Wild Fish e-Harv Wild F | $6-y \mathrm{y}$ <br> Moving Avg | Spawning Year | SpwnrsWild | SpwnrsHatc | Effective <br> Tot. Spwnrs | Wild Fish e-Harv Wild F | 6 -yr <br> Moving Avg |
| 6038 | 0 | 6038 | 8204 |  | 1976 | 5531 | 0 | 5531 | 7515 | 9826 |
| 10322 | 0 | 10322 | 14025 |  | 1977 | 5006 | 0 | 5006 | 6801 | 8807 |
| 8924 | 0 | 8924 | 12125 |  | 1978 | 5968 | 0 | 5968 | 8109 | 8437 |
| 8487 | 0 | 8487 | 11531 |  | 1979 | 7186 | 0 | 7186 | 9764 | 8325 |
| 6447 | 0 | 6447 | 8760 |  | 1980 | 7199 | 0 | 7199 | 9781 | 8267 |
| 3853 | 0 | 3853 | 5235 | 9980 | 1981 | 6140 | 0 | 6140 | 8343 | 8385 |
| 9775 | 0 | 9775 | 13281 | 10826 | 1982 | 5893 | 0 | 5893 | 8006 | 8467 |
| 4686 | 0 | 4686 | 6368 | 9550 | 1983 | 3545 | 0 | 3545 | 4816 | 8136 |
| 8394 | 0 | 8394 | 11405 | 9430 | 1984 | 4221 | 0 | 4221 | 5735 | 7741 |
| 4375 | 0 | 4375 | 5944 | 8499 | 1985 | 7732 | 0 | 7732 | 10505 | 7864 |
| 9394 | 0 | 9394 | 12764 | 9166 | 1986 | 9688 | 0 | 9688 | 13163 | 8428 |
| 8209 | 0 | 8209 | 11154 | 10153 | 1987 | 7501 | 0 | 7501 | 10191 | 8736 |
| 5842 | 0 | 5842 | 7938 | 9262 | 1988 | 8993 | 0 | 8993 | 12219 | 9438 |
| 5862 | 0 | 5862 | 7965 | 9528 | 1989 | 6612 | 0 | 6612 | 8984 | 10133 |
| 5647 | 0 | 5647 | 7673 | 8906 | 1990 | 7854 | 0 | 7854 | 10671 | 10955 |
| 4777 | 0 | 4777 | 6490 | 8997 | 1991 | 3614 | 0 | 3614 | 4910 | 10023 |
| 7115 | 0 | 7115 | 9668 | 8481 | 1992 | 4847 | 0 | 4847 | 6585 | 8927 |
| 5370 | 0 | 5370 | 7296 | 7838 | 1993 | 4017 | 0 | 4017 | 5458 | 8138 |
| 7108 | 0 | 7108 | 9658 | 8125 | 1994 | 3761 | 0 | 3761 | 5110 | 6953 |
| 8714 | 0 | 8714 | 11840 | 8771 | 1995 | 5261 | 0 | 5261 | 7149 | 6647 |
| 9140 | 0 | 9140 | 12419 | 9562 | 1996 | 4503 | 0 | 4503 | 6119 | 5888 |
| 7902 | 0 | 7902 | 10736 | 10269 | 1997 | 5313 | 0 | 5313 | 7219 | 6273 |
| 9074 | 0 | 9074 | 12329 | 10713 | 1998 | 4698 | 0 | 4698 | 6384 | 6240 |
| 7472 | 0 | 7472 | 10153 | 11189 | 1999 | 5829 | 0 | 5829 | 6400 | 6397 |
| 11183 | 0 | 11183 | 15194 | 12112 | 2000 | 6816 | 0 | 6816 | 6885 | 6692 |
| 9504 | 0 | 9504 | 12913 | 12290 |  |  |  |  |  |  |
| 7605 | 0 | 7605 | 10333 | 11943 |  |  |  |  |  |  |
| 7680 | 0 | 7680 | 10435 | 11893 |  |  |  |  |  |  |
| 7456 | 0 | 7456 | 10130 | 11526 |  |  |  |  |  |  |
| 5616 | 0 | 5616 | 7630 | 11106 |  |  |  |  |  |  |


| Basin: Rogue |
| :---: |
| Population: Upper Rogue SR |
| Sub-population: |
| Monitoring sites: Gold Ray Dam |
| Method: Counts made at Gold |
| \|Ray Dam. |
|  |


| Critical Threshold | 258 |
| :---: | :---: |
| Viable Threshold | 897 |
| Last 6-yr Average | 3142 |


Average Distribution of Ages in return year

| Repeat | Age 2 |  | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.10 | 0.01 | 0.15 | 0.54 | 0.19 | 0.00 |  |


| Spawning <br> Year | SpwnrsWild | SpwnrsHatc | Effective Tot. Spwnrs | Harvest Rate | Pre- Hary e-harv Wild Fi |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1974 | 4511 | 723 | 5233 | 0.11 | 5068 |  |
| 1975 | 6573 | 2020 | 8593 | 0.11 | 7385 |  |
| 1976 | 6004 | 1972 | 7976 | 0.11 | 6746 |  |
| 1977 | 2380 | 0 | 2380 | 0.11 | 2674 |  |
| 1978 | 9230 | 445 | 9675 | 0.11 | 10371 |  |
| 1979 | 3542 | 0 | 3542 | 0.11 | 3980 | 6037 |
| 1980 | 10530 | 0 | 10530 | 0.11 | 11831 | 7165 |
| 1981 | 4977 | 0 | 4977 | 0.11 | 5592 | 6866 |
| 1982 | 7080 | 0 | 7080 | 0.11 | 7955 | 7067 |
| 1983 | 8939 | 0 | 8939 | 0.11 | 10044 | 8296 |
| 1984 | 4484 | 0 | 4484 | 0.11 | 5038 | 7407 |
| 1985 | 4543 | 0 | 4543 | 0.11 | 5104 | 7594 |
| 1986 | 7430 | 503 | 7933 | 0.11 | 8348 | 7014 |
| 1987 | 8710 | 723 | 9432 | 0.11 | 9786 | 7713 |
| 1988 | 11534 | 1427 | 12960 | 0.11 | 12959 | 8547 |
| 1989 | 10033 | 148 | 10181 | 0.11 | 11273 | 8751 |
| 1990 | 4996 | 1982 | 6977 | 0.11 | 5613 | 8847 |
| 1991 | 1453 | 2094 | 3547 | 0.11 | 1633 | 8269 |
| 1992 | 2876 | 0 | 2876 | 0.11 | 3231 | 7416 |
| 1993 | 3598 | 645 | 4243 | 0.01 | 3638 | 6391 |
| 1994 | 3620 | 1878 | 5498 | 0.01 | 3660 | 4841 |
| 1995 | 3764 | 2808 | 6572 | 0.01 | 3806 | 3597 |
| 1996 | 5255 | 1165 | 6420 | 0.01 | 5314 | 3547 |
| 1997 | 3127 | 1640 | 4767 | 0.01 | 3161 | 3802 |
| 1998 | 1341 | 175 | 1516 | 0.01 | 1356 | 3489 |
| 1999 | 3087 | 29 | 3116 | 0.01 | 3122 | 3403 |
| 2000 | 2069 | 0 | 2069 | 0.01 | 2092 | 3142 |


| Basin: Rogue |
| :---: |
| Population: Upper Rogue WR |
| Sub-population: |
| Monitoring sites: Gold Ray Dam |
| Method: Counts made at Gold |
| \|Ray Dam. |
|  |


|  | Critical Threshold |
| :--- | ---: |
| Viable Threshold | 235 |
|  | 869 |
|  | 7352 |


Average Distribution of Ages in return year

| Repeat | Age 2 |  | Age 3 | Age 4 | Age 5 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.20 | 0.01 | 0.14 | 0.48 | 0.17 | 0.00 |


| Spawning <br> Year | SpwnrsWild | SpwnrsHatc | Effective <br> Tot. Spwnrs | Harvest <br> Rate | Pre- Harv <br> e-harv Wild Fi | 6 -yr <br> Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1974 | 5570 | 377 | 5947 | 0.08 | 6054 |  |
| 1975 | 6843 | 249 | 7092 | 0.08 | 7438 |  |
| 1976 | 4614 | 0 | 4614 | 0.08 | 5015 |  |
| 1977 | 3800 | 106 | 3905 | 0.08 | 4130 |  |
| 1978 | 4512 | 0 | 4512 | 0.08 | 4904 |  |
| 1979 | 8980 | 912 | 9892 | 0.08 | 9761 | 6217 |
| 1980 | 8156 | 1194 | 9349 | 0.08 | 8865 | 6686 |
| 1981 | 5271 | 617 | 5887 | 0.08 | 5729 | 6401 |
| 1982 | 4213 | 442 | 4655 | 0.08 | 4579 | 6328 |
| 1983 | 6573 | 0 | 6573 | 0.08 | 7145 | 6831 |
| 1984 | 5009 | 0 | 5009 | 0.08 | 5445 | 6921 |
| 1985 | 8255 | 0 | 8255 | 0.08 | 8973 | 6789 |
| 1986 | 10643 | 554 | 11197 | 0.08 | 11569 | 7240 |
| 1987 | 11663 | 854 | 12517 | 0.08 | 12677 | 8398 |
| 1988 | 10103 | 436 | 10539 | 0.08 | 10982 | 9465 |
| 1989 | 8675 | 1028 | 9703 | 0.08 | 9429 | 9846 |
| 1990 | 6183 | 343 | 6526 | 0.08 | 6721 | 10059 |
| 1991 | 2685 | 417 | 3103 | 0.08 | 2919 | 9050 |
| 1992 | 2955 | 174 | 3129 | 0.08 | 3212 | 7657 |
| 1993 | 4310 | 401 | 4711 | 0.01 | 4345 | 6268 |
| 1994 | 4900 | 174 | 5074 | 0.01 | 4940 | 5261 |
| 1995 | 8559 | 662 | 9220 | 0.01 | 8628 | 5128 |
| 1996 | 7279 | 0 | 7279 | 0.01 | 7338 | 5230 |
| 1997 | 11426 | 341 | 11767 | 0.01 | 11518 | 6664 |
| 1998 | 3744 | 0 | 3744 | 0.01 | 3774 | 6757 |
| 1999 | 7665 | 0 | 7665 | 0.01 | 7727 | 7321 |
| 2000 | 5087 | 0 | 5087 | 0.01 | 5128 | 7352 |
|  |  |  |  |  |  |  |


| Basin: Rogue |
| :---: |
| Population: Middle Rogue SR |
| Sub-population: |
| Monitoring sites: Foots Cr |
| Kane $\mathbf{C r}$ |
|  |
| Method: Redd Surveys |


|  |  |
| :--- | :---: |
| Critical Threshold | 5.30 |
|  |  |
| Viable Threshold | 9.42 |
|  |  |
|  |  |


Average Distribution of Ages at time of spawning

| Repeat | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |


| Spawning Year | SpwnrsWild | SpwnrsHatc | Effective Tot. Spwnrs | Wild Fish Harv Rate | Pre- Harv <br> e-harv Wild Fi | $6-\mathrm{yr}$ Moving Avg |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1977 | 72.4 | 0 | 72.4 | 0.08 | 78.7 |  |
| 1978 | 106.8 | 0 | 106.8 | 0.08 | 116.1 |  |
| 1979 | 112.9 | 0 | 112.9 | 0.08 | 122.8 |  |
| 1980 | 204.3 | 0 | 204.3 | 0.08 | 222.1 |  |
| 1981 | 94.0 | 0 | 94.0 | 0.08 | 102.2 |  |
| 1982 | 18.2 | 0 | 18.2 | 0.08 | 19.8 | 110.3 |
| 1983 | 51.0 | 0 | 51.0 | 0.08 | 55.4 | 106.4 |
| 1984 | 48.8 | 0 | 48.8 | 0.08 | 53.1 | 95.9 |
| 1985 | 43.4 | 0 | 43.4 | 0.08 | 47.1 | 83.3 |
| 1986 | 53.6 | 0 | 53.6 | 0.08 | 58.3 | 56.0 |
| 1987 | 68.2 | 0 | 68.2 | 0.08 | 74.2 | 51.3 |
| 1988 | 22.0 | 0 | 22.0 | 0.08 | 24.0 | 52.0 |
| 1989 | 56.7 | 0 | 56.7 | 0.08 | 61.6 | 53.0 |
| 1990 | 121.3 | 0 | 121.3 | 0.08 | 131.9 | 66.2 |
| 1991 | 15.4 | 0 | 15.4 | 0.08 | 16.7 | 61.1 |
| 1992 | 2.8 | 0 | 2.8 | 0.08 | 3.0 | 51.9 |
| 1993 | 6.5 | 0 | 6.5 | 0.08 | 7.1 | 40.7 |
| 1994 | 0.9 | 0 | 0.9 | 0.08 | 1.0 | 36.9 |
| 1995 | 7.7 | 0 | 7.7 | 0.08 | 8.4 | 28.0 |
| 1996 | 5.8 | 0 | 5.8 | 0.01 | 5.8 | 7.0 |
| 1997 | 20.7 | 0 | 20.7 | 0.01 | 20.8 | 7.7 |
| 1998 | 10.0 | 0 | 10.0 | 0.01 | 10.1 | 8.9 |
| 1999 | 15.4 | 0 | 15.4 | 0.01 | 15.5 | 10.3 |
| 2000 | 44.7 | 0 | 44.7 | 0.01 | 45.0 | 17.6 |


| Basin: Rogue |
| :---: |
| Population: Applegate WR |
| Sub-population: |
| Monitoring sites: Applgate Dam Trap |
| Method: Trap counts and |
| \|ratio of hatchery and |
| wild fish in downstream |
| fishery. |


| Critical Threshold | 59 |
| :--- | :---: |
|  |  |
| Viable Threshold | 210 |
|  | 1371 |



| Spawning <br> Year |  |  | SpwnrsWild | SpwnrsHatc | Effective <br> Tot. Spwnrs | Harvest <br> Rate |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1983 | 273 | 91 | 364 | 0.19 | Pre-Harv <br> e-harv Wild Fi | 6-yr <br> Moving Avg |
| 1984 | 733 | 245 | 979 | 0.19 | 906 |  |
| 1985 | 621 | 208 | 829 | 0.19 | 767 |  |
| 1986 | 286 | 96 | 382 | 0.19 | 353 |  |
| 1987 | 624 | 209 | 832 | 0.19 | 770 |  |
| 1988 | 454 | 152 | 606 | 0.19 | 561 | 616 |
| 1989 | 1891 | 632 | 2523 | 0.19 | 2335 | 949 |
| 1990 | 565 | 189 | 754 | 0.19 | 697 | 914 |
| 1991 | 359 | 120 | 478 | 0.19 | 443 | 860 |
| 1992 | 229 | 76 | 305 | 0.19 | 282 | 848 |
| 1993 | 540 | 148 | 687 | 0.01 | 545 | 810 |
| 1994 | 188 | 51 | 239 | 0.01 | 190 | 749 |
| 1995 | 1490 | 407 | 1898 | 0.01 | 1505 | 610 |
| 1996 | 1153 | 315 | 1468 | 0.01 | 1164 | 688 |
| 1997 | 1453 | 397 | 1851 | 0.01 | 1467 | 859 |
| 1998 | 1391 | 380 | 1771 | 0.01 | 1404 | 1046 |
| 1999 | 1458 | 399 | 1857 | 0.01 | 1472 | 1200 |
| 2000 | 1203 | 329 | 1532 | 0.01 | 1214 | 1371 |

Appendix 3. Summary of productivity estimates (Ricker $a$-values), associated regression statistics $\left(\mathrm{R}^{2}\right)$, and proportion of hatchery fish $\left(\mathrm{P}_{\mathrm{h}}\right)$ for 27 populations of Oregon steelhead by 7 -year moving sequences of spawner and recruit data, 1974-95 brood years.

| Sequence <br> Years | Rogue Summers |  |  |  | MidRogue <br> Summers |  |  | Rogue Winters |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |  |
| $1974-80$ | 1.290 | 0.96 | 0.10 |  |  |  | 0.717 | 0.81 | 0.05 |  |
| $1975-81$ | 1.191 | 0.90 | 0.08 |  |  |  | 0.822 | 0.83 | 0.06 |  |
| $1976-82$ | 1.181 | 0.90 | 0.04 |  |  |  | 1.005 | 0.77 | 0.06 |  |
| $1977-83$ | 1.149 | 0.86 | 0.01 | 0.805 | 0.81 | 0.00 | 1.014 | 0.66 | 0.06 |  |
| $1978-84$ | 1.241 | 0.79 | 0.01 | 0.565 | 0.79 | 0.00 | 1.323 | 0.67 | 0.06 |  |
| $1979-85$ | 1.332 | 0.79 | 0.00 | 0.655 | 0.82 | 0.00 | 1.743 | 0.95 | 0.06 |  |
| $1980-86$ | 1.360 | 0.73 | 0.01 | 0.737 | 0.81 | 0.00 | 1.603 | 0.95 | 0.05 |  |
| $1981-87$ | 1.819 | 0.69 | 0.02 | 1.300 | 0.75 | 0.00 | 1.741 | 0.96 | 0.05 |  |
| $1982-88$ | 1.913 | 0.84 | 0.04 | 0.302 | 0.05 | 0.00 | 1.927 | 0.93 | 0.04 |  |
| $1983-89$ | 1.930 | 0.85 | 0.04 | -0.529 | 0.00 | 0.00 | 2.002 | 0.89 | 0.04 |  |
| $1984-90$ | 1.723 | 0.87 | 0.08 | 0.782 | 0.46 | 0.00 | 1.517 | 0.83 | 0.05 |  |
| $1985-91$ | 1.071 | 0.83 | 0.16 | 0.136 | 0.38 | 0.00 | 1.306 | 0.87 | 0.06 |  |
| $1986-92$ | 0.772 | 0.92 | 0.16 | 0.302 | 0.54 | 0.00 | 1.353 | 0.94 | 0.07 |  |
| $1987-93$ | 0.615 | 0.92 | 0.18 | 0.319 | 0.73 | 0.00 | 1.494 | 0.97 | 0.08 |  |
| $1988-94$ | 0.433 | 0.80 | 0.21 | 0.801 | 0.74 | 0.00 | 1.560 | 0.98 | 0.07 |  |
| $1989-95$ | 0.536 | 0.69 | 0.26 | 1.062 | 0.80 | 0.00 | 1.417 | 0.92 | 0.08 |  |


| Sequence <br> Years | Applegate |  |  |  |  | N. Umpqua <br> Summers |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | a | $\mathrm{R}^{2}$ | P . Umpqua Winters |  |  |  |
| $1974-80$ |  |  |  | 1.237 | 0.49 | 0.08 | 1.726 | 0.66 | 0.00 |
| $1975-81$ |  |  |  | 1.001 | 0.18 | 0.07 | 2.177 | 0.65 | 0.00 |
| $1976-82$ |  |  |  | 1.121 | 0.18 | 0.07 | 2.277 | 0.56 | 0.00 |
| $1977-83$ |  |  |  | 1.268 | 0.27 | 0.06 | 2.133 | 0.73 | 0.00 |
| $1978-84$ |  |  |  | 1.192 | 0.20 | 0.05 | 2.278 | 0.80 | 0.00 |
| $1979-85$ |  |  |  | 1.631 | 0.73 | 0.05 | 2.097 | 0.79 | 0.00 |
| $1980-86$ |  |  |  | 1.904 | 0.82 | 0.05 | 1.725 | 0.91 | 0.00 |
| $1981-87$ |  |  |  | 2.002 | 0.87 | 0.05 | 1.781 | 0.80 | 0.00 |
| $1982-88$ |  |  |  | 1.835 | 0.84 | 0.07 | 1.828 | 0.83 | 0.00 |
| $1983-89$ | 0.367 | 0.60 | 0.25 | 1.365 | 0.76 | 0.09 | 1.641 | 0.75 | 0.00 |
| $1984-90$ | 0.242 | 0.56 | 0.25 | 0.926 | 0.74 | 0.10 | 1.384 | 0.57 | 0.00 |
| $1985-91$ | 0.484 | 0.55 | 0.25 | 0.718 | 0.77 | 0.12 | 0.811 | 0.54 | 0.00 |
| $1986-92$ | 0.670 | 0.60 | 0.25 | 0.923 | 0.88 | 0.12 | 0.820 | 0.69 | 0.00 |
| $1987-93$ | 0.844 | 0.59 | 0.25 | 1.064 | 0.93 | 0.12 | 1.179 | 0.94 | 0.00 |
| $1988-94$ | 1.251 | 0.70 | 0.24 | 1.359 | 0.96 | 0.11 | 1.152 | 0.97 | 0.00 |
| $1989-95$ | 1.487 | 0.86 | 0.24 | 1.364 | 0.86 | 0.09 |  |  |  |

Appendix 3. (Continued)

| Sequence <br> Years | Salmonberry |  |  | Calapooia |  |  |  | Lower S. Santiam |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\mathbf{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |  |
| $1974-80$ | 0.124 | 0.30 | 0.00 |  |  |  |  |  |  |  |
| $1975-81$ | 0.028 | 0.25 | 0.00 |  |  |  |  |  |  |  |
| $1976-82$ | 1.557 | 0.51 | 0.00 |  |  |  |  |  |  |  |
| $1977-83$ | 2.069 | 0.81 | 0.00 |  |  |  |  |  |  |  |
| $1978-84$ | 1.983 | 0.82 | 0.00 |  |  |  |  |  |  |  |
| $1979-85$ | 1.715 | 0.73 | 0.00 |  |  |  |  |  |  |  |
| $1980-86$ | 1.825 | 0.91 | 0.00 | 1.912 | 0.86 | 0.00 |  |  |  |  |
| $1981-87$ | 1.776 | 0.94 | 0.00 | 2.033 | 0.87 | 0.00 |  |  |  |  |
| $1982-88$ | 1.695 | 0.93 | 0.00 | 1.262 | 0.25 | 0.00 |  |  |  |  |
| $1983-89$ | 1.573 | 0.78 | 0.00 | -0.142 | 0.04 | 0.00 | 0.197 | 0.17 | 0.48 |  |
| $1984-90$ | 1.115 | 0.76 | 0.00 | -0.317 | 0.06 | 0.00 | 0.322 | 0.21 | 0.44 |  |
| $1985-91$ | 0.839 | 0.73 | 0.00 | -0.643 | 0.03 | 0.00 | 0.175 | 0.06 | 0.35 |  |
| $1986-92$ | 0.776 | 0.71 | 0.00 | -0.898 | 0.01 | 0.00 | -1.232 | 0.01 | 0.26 |  |
| $1987-93$ | 0.347 | 0.66 | 0.00 | 0.064 | 0.24 | 0.00 | -0.438 | 0.01 | 0.17 |  |
| $1988-94$ | 0.054 | 0.21 | 0.00 | 0.781 | 0.49 | 0.00 | -0.064 | 0.08 | 0.09 |  |
| $1989-95$ |  |  |  | 1.034 | 0.44 | 0.00 | -0.182 | 0.20 | 0.04 |  |
| $1990-96$ |  |  |  | 1.277 | 0.60 | 0.00 | 0.451 | 0.52 | 0.01 |  |


| Sequence <br> Years | Upper S. Santiam |  |  | N. Santiam |  |  | Molalla |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |  |
| $1974-80$ | 0.400 | 0.22 | 0.00 |  |  |  |  |  |  |  |
| $1975-81$ | 0.645 | 0.28 | 0.00 |  |  |  |  |  |  |  |
| $1976-82$ | 0.776 | 0.41 | 0.06 |  |  |  |  |  |  |  |
| $1977-83$ | 0.606 | 0.53 | 0.11 |  |  |  |  |  |  |  |
| $1978-84$ | 0.307 | 0.64 | 0.20 |  |  |  |  |  |  |  |
| $1979-85$ | 0.427 | 0.67 | 0.29 |  |  |  |  |  |  |  |
| $1980-86$ | 0.406 | 0.60 | 0.38 |  |  |  | 0.735 | 0.75 | 0.46 |  |
| $1981-87$ | 0.122 | 0.43 | 0.46 |  |  |  | 0.473 | 0.29 | 0.46 |  |
| $1982-88$ | -0.052 | 0.41 | 0.51 |  |  |  | 0.701 | 0.45 | 0.46 |  |
| $1983-89$ | -0.110 | 0.42 | 0.48 | -0.146 | 0.06 | 0.15 | 0.010 | 0.11 | 0.46 |  |
| $1984-90$ | -0.376 | 0.35 | 0.44 | -0.384 | 0.02 | 0.15 | -0.704 | 0.01 | 0.46 |  |
| $1985-91$ | 0.340 | 0.77 | 0.35 | -1.121 | 0.06 | 0.15 | -1.278 | 0.02 | 0.46 |  |
| $1986-92$ | 0.159 | 0.57 | 0.26 | -1.192 | 0.03 | 0.15 | -1.808 | 0.11 | 0.43 |  |
| $1987-93$ | 0.268 | 0.55 | 0.17 | -0.416 | 0.08 | 0.15 | -0.593 | 0.14 | 0.40 |  |
| $1988-94$ | 0.418 | 0.61 | 0.09 | 0.030 | 0.20 | 0.15 | -0.513 | 0.18 | 0.36 |  |
| $1989-95$ | 1.805 | 0.84 | 0.04 | 0.984 | 0.56 | 0.14 | 0.040 | 0.42 | 0.33 |  |
| $1990-96$ | 1.745 | 0.89 | 0.01 | 1.328 | 0.68 | 0.14 | 0.606 | 0.47 | 0.30 |  |

Appendix 3. (Continued)

| Sequence <br> Years | Clackamas |  |  | Sandy |  |  | Warm Springs |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |
|  | 1.559 | 0.84 | 0.21 |  |  |  |  |  |  |
| $1975-81$ | 1.465 | 0.81 | 0.26 |  |  |  |  |  |  |
| $1976-82$ | 1.461 | 0.82 | 0.28 |  |  |  |  |  |  |
| $1977-83$ | 1.262 | 0.85 | 0.27 |  |  |  |  |  |  |
| $1978-84$ | 1.306 | 0.89 | 0.26 | 0.439 | 0.53 | 0.50 |  |  |  |
| $1979-85$ | 1.181 | 0.83 | 0.22 | 0.308 | 0.33 | 0.47 |  |  |  |
| $1980-86$ | 1.114 | 0.75 | 0.19 | 0.568 | 0.47 | 0.46 | 1.630 | 0.70 | 0.00 |
| $1981-87$ | 0.891 | 0.46 | 0.18 | 1.302 | 0.46 | 0.49 | 1.388 | 0.44 | 0.00 |
| $1982-88$ | 1.411 | 0.52 | 0.16 | 0.534 |  | 0.48 | 1.549 | 0.71 | 0.00 |
| $1983-89$ | 1.257 | 0.47 | 0.16 | -0.615 |  | 0.50 | 1.311 | 0.55 | 0.00 |
| $1984-90$ | 0.969 | 0.37 | 0.20 | -1.142 |  | 0.49 | 0.911 | 0.48 | 0.00 |
| $1985-91$ | 0.221 | 0.35 | 0.24 | -0.602 | 0.03 | 0.51 | -0.232 | 0.15 | 0.00 |
| $1986-92$ | 1.049 | 0.41 | 0.26 | -1.105 | 0.01 | 0.54 | -0.102 | 0.41 | 0.00 |
| $1987-93$ | 0.587 | 0.27 | 0.26 | -0.760 | 0.18 | 0.56 | 0.052 | 0.58 | 0.00 |
| $1988-94$ | 0.078 | 0.16 | 0.25 | -0.536 | 0.37 | 0.55 |  |  |  |
| $1989-95$ | 0.636 | 0.36 | 0.26 | -0.304 | 0.53 | 0.55 |  |  |  |


| Sequence <br> Years | Deschutes |  |  |  | Lower NFk. John <br> Day |  |  |  | Upper NFk John <br> Day |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $a$ | $R^{2}$ | $P_{h}$ | $a$ | $R^{2}$ | $P_{h}$ | $a$ | $R^{2}$ | $P_{h}$ |  |  |
| $1974-80$ |  |  |  |  |  |  |  |  |  |  |  |
| $1975-81$ |  |  |  |  |  |  |  |  |  |  |  |
| $1976-82$ |  |  |  | 3.162 | 0.70 | 0.00 |  |  |  |  |  |
| $1977-83$ |  |  |  | 2.896 | 0.69 | 0.00 | 1.207 | 0.14 | 0.00 |  |  |
| $1978-84$ | 1.464 | 0.94 | 0.33 | 2.368 | 0.46 | 0.00 | 1.631 | 0.35 | 0.00 |  |  |
| $1979-85$ | 1.511 | 0.96 | 0.32 | 2.466 | 0.80 | 0.00 | 2.282 | 0.74 | 0.00 |  |  |
| $1980-86$ | 1.377 | 0.94 | 0.32 | 2.091 | 0.84 | 0.00 | 2.676 | 0.93 | 0.00 |  |  |
| $1981-87$ | 1.300 | 0.93 | 0.35 | 2.016 | 0.88 | 0.00 | 2.796 | 0.94 | 0.00 |  |  |
| $1982-88$ | 1.405 | 0.78 | 0.38 | 1.365 | 0.80 | 0.00 | 2.152 | 0.76 | 0.00 |  |  |
| $1983-89$ | -0.993 | 0.04 | 0.41 | 0.841 | 0.82 | 0.00 | 1.439 | 0.59 | 0.00 |  |  |
| $1984-90$ | -1.839 | 0.06 | 0.43 | 0.511 | 0.95 | 0.00 | 2.049 | 0.73 | 0.00 |  |  |
| $1985-91$ | -1.285 | 0.04 | 0.42 | 0.489 | 0.95 | 0.00 | 2.701 | 0.75 | 0.00 |  |  |
| $1986-92$ | -1.204 | 0.10 | 0.44 | 0.444 | 0.92 | 0.00 | 2.687 | 0.75 | 0.00 |  |  |
| $1987-93$ | -0.393 | 0.48 | 0.50 | 0.709 | 0.74 | 0.00 | 2.351 | 0.61 | 0.00 |  |  |
| $1988-94$ |  |  |  | 1.407 | 0.82 | 0.00 | 2.736 | 0.64 | 0.00 |  |  |
| $1989-95$ |  |  |  |  |  |  |  |  |  |  |  |

Appendix 3. (Continued)

| Sequence <br> Years | Middle Fk. John <br> Day |  |  |  | South Fk. John Day |  |  | Lower John Day |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $\boldsymbol{a}$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |  |
| $1974-80$ | 1.171 | 0.83 | 0.00 | 1.663 | 0.94 | 0.00 | 3.470 | 0.90 | 0.00 |  |
| $1975-81$ | 1.328 | 0.73 | 0.00 | 1.809 | 0.92 | 0.00 | 3.581 | 0.90 | 0.00 |  |
| $1976-82$ | 1.621 | 0.78 | 0.00 | 2.000 | 0.76 | 0.00 | 3.816 | 0.91 | 0.00 |  |
| $1977-83$ | 1.832 | 0.79 | 0.00 | 1.713 | 0.65 | 0.00 | 3.878 | 0.90 | 0.00 |  |
| $1978-84$ | 2.050 | 0.73 | 0.00 | 1.391 | 0.46 | 0.00 | 3.635 | 0.86 | 0.00 |  |
| $1979-85$ | 1.888 | 0.62 | 0.00 | 1.930 | 0.79 | 0.00 | 3.467 | 0.97 | 0.00 |  |
| $1980-86$ | 2.053 | 0.90 | 0.00 | 2.258 | 0.81 | 0.00 | 2.607 | 0.83 | 0.00 |  |
| $1981-87$ | 2.193 | 0.94 | 0.00 | 2.416 | 0.89 | 0.00 | 2.076 | 0.81 | 0.00 |  |
| $1982-88$ | 2.129 | 0.94 | 0.00 | 2.431 | 0.88 | 0.00 | 2.110 | 0.84 | 0.00 |  |
| $1983-89$ | 1.825 | 0.80 | 0.00 | 1.289 | 0.75 | 0.00 | 1.142 | 0.62 | 0.00 |  |
| $1984-90$ | 1.066 | 0.63 | 0.00 | 0.842 | 0.79 | 0.00 | -0.054 | 0.36 | 0.00 |  |
| $1985-91$ | -0.090 | 0.12 | 0.00 | 0.313 | 0.82 | 0.00 | -0.721 | 0.39 | 0.00 |  |
| $1986-92$ | -0.237 | 0.08 | 0.00 | 0.094 | 0.66 | 0.00 | -0.702 | 0.51 | 0.00 |  |
| $1987-93$ | -0.220 | 0.10 | 0.00 | -0.089 | 0.57 | 0.00 | -0.574 | 0.59 | 0.00 |  |
| $1988-94$ | -0.111 | 0.24 | 0.00 | -0.247 | 0.29 | 0.00 | 0.085 | 0.41 | 0.00 |  |
| $1989-95$ |  |  |  |  |  |  |  |  |  |  |


| Sequence <br> Years | Upper John Day |  |  |  | Umatilla |  |  | Upper Grande <br> Ronde |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | $a$ | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |  |
| $1974-80$ | 1.858 | 0.86 | 0.00 | 0.606 | 0.15 | 0.00 | 1.419 | 0.04 | 0.00 |  |
| $1975-81$ | 2.239 | 0.86 | 0.00 | 0.444 | 0.09 | 0.00 | 2.447 | 0.21 | 0.00 |  |
| $1976-82$ | 2.394 | 0.77 | 0.00 | 1.384 | 0.45 | 0.00 | 2.946 | 0.26 | 0.00 |  |
| $1977-83$ | 2.436 | 0.72 | 0.00 | 1.930 | 0.68 | 0.00 | 3.579 | 0.64 | 0.00 |  |
| $1978-84$ | 2.222 | 0.45 | 0.00 | 1.930 | 0.68 | 0.00 | 3.937 | 0.90 | 0.00 |  |
| $1979-85$ | 2.407 | 0.80 | 0.00 | 2.333 | 0.96 | 0.00 | 2.824 | 0.64 | 0.00 |  |
| $1980-86$ | 2.025 | 0.84 | 0.00 | 2.244 | 0.96 | 0.00 | 2.325 | 0.55 | 0.00 |  |
| $1981-87$ | 2.117 | 0.91 | 0.00 | 2.327 | 0.94 | 0.00 | 2.187 | 0.53 | 0.00 |  |
| $1982-88$ | 2.017 | 0.91 | 0.00 | 2.270 | 0.98 | 0.00 | 1.327 | 0.48 | 0.02 |  |
| $1983-89$ | 1.362 | 0.74 | 0.00 | 2.200 | 0.94 | 0.01 | 0.148 | 0.80 | 0.03 |  |
| $1984-90$ | 0.691 | 0.57 | 0.00 | 1.663 | 0.75 | 0.03 | 0.028 | 0.88 | 0.05 |  |
| $1985-91$ | -0.301 | 0.37 | 0.00 | -0.024 | 0.15 | 0.06 | 0.151 | 0.84 | 0.08 |  |
| $1986-92$ | -0.594 | 0.11 | 0.00 | 0.139 | 0.60 | 0.11 | 0.735 | 0.66 | 0.12 |  |
| $1987-93$ | -0.475 | 0.16 | 0.00 | 0.213 | 0.63 | 0.14 | 0.718 | 0.65 | 0.15 |  |
| $1988-94$ | -0.473 | 0.17 | 0.00 | 0.114 | 0.52 | 0.19 |  |  |  |  |

## Appendix 3.(Continued)

| Sequence <br> Years | Lower Grande <br> Ronde |  |  |  | Joseph |  |  | Imnaha |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | a | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ | a | $\mathrm{R}^{2}$ | $\mathrm{P}_{\mathrm{h}}$ |  |
| $1974-80$ | 1.178 | 3.7 | 0.00 | 1.966 | 0.26 | 0.00 | 1.478 | 0.46 | 0.00 |  |
| $1975-81$ | 1.245 | 3.6 | 0.00 | 1.968 | 0.16 | 0.00 | 1.644 | 0.44 | 0.00 |  |
| $1976-82$ | 1.425 | 3.3 | 0.00 | 2.025 | 0.17 | 0.00 | 1.735 | 0.38 | 0.00 |  |
| $1977-83$ | 1.593 | 3.1 | 0.00 | 2.282 | 0.23 | 0.00 | 1.583 | 0.21 | 0.00 |  |
| $1978-84$ | 1.724 | 2.9 | 0.00 | 2.716 | 0.52 | 0.00 | 1.995 | 0.39 | 0.00 |  |
| $1979-85$ | 1.648 | 3.3 | 0.00 | 2.588 | 0.87 | 0.00 | 2.176 | 0.66 | 0.00 |  |
| $1980-86$ | 1.397 | 4.1 | 0.00 | 2.300 | 0.98 | 0.00 | 2.565 | 0.92 | 0.00 |  |
| $1981-87$ | 1.441 | 3.7 | 0.00 | 2.312 | 0.98 | 0.00 | 2.440 | 0.94 | 0.03 |  |
| $1982-88$ | 1.026 | 4.6 | 0.02 | 2.296 | 0.98 | 0.00 | 1.859 | 0.75 | 0.06 |  |
| $1983-89$ | 0.486 | 7.7 | 0.04 | 2.359 | 0.96 | 0.00 | 1.466 | 0.69 | 0.09 |  |
| $1984-90$ | 0.058 | 17.4 | 0.06 | 2.130 | 0.87 | 0.00 | 1.188 | 0.66 | 0.11 |  |
| $1985-91$ | -0.018 | 18.0 | 0.09 | 1.082 | 0.79 | 0.00 | 0.195 | 0.42 | 0.14 |  |
| $1986-92$ | 0.034 | 10.1 | 0.11 | 1.172 | 0.89 | 0.00 | 0.301 | 0.63 | 0.17 |  |
| $1987-93$ | 0.191 | 7.2 | 0.13 | 1.196 | 0.87 | 0.00 | 0.336 | 0.65 | 0.20 |  |
| $1988-94$ |  |  |  |  |  |  | 0.275 | 0.62 | 0.20 |  |
| $1989-95$ |  |  |  |  |  |  |  |  |  |  |

