# POPULATION DYNAMICS OF WHITE BASS AND STRIPED BASS X WHITE BASS HYBRIDS <br> IN LAKE CARL BLACKWELL, OKLAHOMA 

By<br>MAURICE IBECHEMA MUONEKE<br>Bachelor of Science Nicholls State University Thibodaux, Louisiana 1979<br>Master of Science Louisiana State University Baton Rouge, Louisiana 1982

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IN LAKE CARL BLACKWELL, OKLAHOMA

Thesis Approved:


## PREFACE

Striped bass Morone saxatilis $x$ white bass M. chrysops hybrids are stocked in numerous reservoirs in Oklahoma, some of which contain populations of striped bass and (or) white bass. There is some uncertainty in the use of univariate morphometric characters, developed for use in the southeastern states, for separating the hybrid from the parental species in oklahoma. A multivariate approach employing discriminant analysis and sheared principal component analysis was used to identify characters for separation of the three groups. A conceptual simulation model was developed for the white bass and striped bass $x$ white bass fisheries in reservoirs.

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## CHAPTER I

## INTRODUCTION

## Statement of Problems

The demand for a pelagic sport fish capable of utilizing clupeid forage fish abundant in numerous reservoirs has enhanced the popularity of striped bass, Morone saxatilis $x$ white bass M. chrysops hybrid fisheries in many southeastern States, including Oklahoma. In many cases, this demand has resulted in striped bass, Morone saxatilis $x$ white bass, M. chrysops hybrids being stocked into reservoirs with populations of striped bass and (or) white bass. There is concern that the hybrid, by virtue of its voracious appetite, fast growth rate, and adaptability to a wide range of habitat types, may have adverse impacts on other sport fishes such as white bass, walleye, crappie, and largemouth bass.

Since the introduction of striped bass $x$ white bass hybrids into Oklahoma reservoirs in the late seventies, the hybrid has grown in popularity to the extent that some oklahoma reservoirs are designated as "hybrid" reservoirs. Often these "hybrid" reservoirs also contain white bass or striped bass. In recent years there have been some problems
with the use of morphometric and meristic characters that were developed for separating oklahoma hybrids from the parental species. Univariate characters have proven to be inadequate for unambiguous separation of Morone sp. in Oklahoma reservoirs. The ability to identify accurately is necessary because of the importance of the Morone sp. to sport fishery in Oklahoma.

Fish populations are controlled by habitat limitations, density-dependent factors, and stochastic environmental variations. Most fishery management strategies are formulated to account for variations in population sizes due to density-dependent and habitat related causes but exclude variations due to stochastic events. White bass and striped bass $x$ striped bass hybrids in Lake Carl Blackwell (Maughan et al. 1986) do not appear to be limited by habitat but may be limited by stochastic environmental factors. They also may be limited by density dependent factors because white bass and hybrids have similar ecological requirements, often occur in the same reservoirs and are often managed as one fishery. The popularity of these fish makes it important to understand the population dynamics and ecology of the two groups and their effects on other native fish populations.

## Objectives

The inability to separate striped bass $x$ white bass hybrids from both parental forms, with univariate charac-
ters, developed for use in the southeastern states, created a need for identifying groups of characters that could be used for separating the three forms. Therefore, I developed multivariate criteria for separation of the three groups. Fifty-two morphometric characters were measured on each fish with the objective of using linear discriminant function analysis and sheared principal component analysis to elucidate the morphological differences between the three groups. The goal of the analysis was to identify 'groups' of morphometric variables that would facilitate separation.

The second objective of this study was to use as much of the available information as possible to develop a conceptual model of the population dynamics of striped bass $x$ white bass hybrids and white bass in Lake Carl Blackwell, Oklahoma. The major assumptions used in the development of this model were that the hybrid and white bass populations were constrained by density-dependent mechanisms and subject to stochastic environmental variations. The model utilizes information routinely collected in fish population surveys and if the underlying assumptions are correct, can be used to evaluate the effects of various management practices on harvest, yield, population size, and population structure.

## Dissertation Format

This dissertation consists of three chapters. The last two chapters have been prepared as independent documents to
be submitted for publication. References, tables, and figures for chapters II and III are included jointly at the end of the dissertation.

## CHAPTER II

# COMPARATIVE MORPHOLOGY OE THE WHITE BASS AND STRIPED BASS $X$ WHITE BASS HYBRIDS IN LAKE CARL BLACKWELL, OKLAHOMA 

## Literature Review

Hybrids of striped bass Morone saxatilis and white bass M. chrysops crosses are of considerable interest to taxonomy, phylogeny, and management. Striped bass $x$ white bass hybrids have been stocked into several reservoir/river systems which contain endemic white bass populations (Ware 1975) and there is concern that the hybrid may be reproducing or backcrossing with either or both of the parental species within those aquatic systems (Avise and van Den Avyle 1984; Crawford et al. 1984).

The introduction of striped bass $x$ white bass hybrids (hereafter referred to as the hybrid) into reservoirs has also created questions of identification among fishermen, fishery managers, game rangers, and even fishery biologists. The commonly accepted criterion for separation of the hybrid from white bass is the shape of the tooth patches on the
base of the tongue. The hybrid and striped bass are sup-
posed to have split patches whereas the white bass has only one patch (Williams 1972). This character, however, does not accurately separate the three groups in Oklahoma. It is possible that the problem of identifying the hybrid in oklahoma waters may be exacerbated by the fact that parental brood stock as well as fingerlings have come from locations in South Carolina, Louisiana, Illinois, and locally from Oklahoma (Harold Namminga: Oklahoma Department of Wildife Conservation (ODWC)). This diversity of parental stock may have resulted in a diverse gene pool and high levels of phenotypic variability.

Morphometric and meristic characters of Morone sp. hybrids in the southeastern U.S. have been described in some detail (Bayless 1968, 1972; Harrell 1984; Kerby 1979a, 1979b; Kerby et al. 1971; Williams 1976). Harrell (1984) successfully used osteological, morphometric, and meristic characters to identify Morone hybrids. Kerby (1979b) compared the morphometric characters of the striped bass $x$ white bass hybrid with those of the parental species and established some ratios which have also been used to separate the hybrid from the white bass. Kerby (1979b) found a linear relationship between most body length measurements and standard length in the striped bass $x$ white bass hybrid as well as in the striped bass $x$ white perch hybrid. Based partly on Kerby et al.'s (1971) data, Williams (1976) concluded that the ratios of body depth/head length and fork length/body depth could be used to separate striped bass from either white bass or striped
bass $x$ white bass hybrids. These ratios (Williams 1976) (fork length/body depth for white bass, striped bass, and the hybrid were 3.477 , 4.440 , and 3.460 respectively and the corresponding body depth/head length ratios were 1.198 , 0.893, and 1.158 ) have been widely used to identify these forms. All these authors are in agreement that no single character always separates the three forms from one another. Also many of the ratios and relationships used elsewhere do not seem to be applicable to Oklahoma fish.

Since no single character can best be described as "definitive" in its power to separate the white bass from the striped bass $x$ white bass hybrid $I$ have proposed development of a multivariate approach. The multivariate techniques $I$ have used in this development are the linear discriminant analysis, principal component analysis and a modified (sheared) principal component analysis.

## Materials and Methods

## Sample Collection and Preparation

Fish collections (gill net) were made from April to November 1985 by biologists of the Oklahoma Department of Wildife Conservation at Fort Supply Lake, Lake Texoma, Optima Lake, Canton Lake, and the Great Salt Plains Lake. I also made gill net collections from Lake Carl Blackwell during the same period.

Known striped bass $x$ white bass hybrids were obtained
from the Southeastern Fish Culture Laboratory, United States Fish and Wildlife Service, Marion, Alabama and known striped bass were obtained from collections made in Lake Texoma, Oklahoma, by personnel of the ODWC. White bass were collected from Lake Carl Blackwell and their identity was electrophoretically verified by the ODWC's Fishery Research Laboratory. The electrophoretic identification of the white bass was done using the starch gel methods of Selander et al. (1971). The interpretation of the results follows those reported by Avise and Van Den Avyle (1984). The three enzyme loci used to confirm identification were: (i) phosphoglucoisomerase (PGI), (ii) calcium binding protein (CBP), and (iii) esterase (EST).

Upon capture, fish suitable for morphometric and meristic analyses were immediately packed on ice and the livers, to be used for electrophoresis, were immediately removed and frozen. The fish were then injected with $10 \%$ formalin (to prevent visceral deterioration) and subsequently preserved in $10 \%$ formalin.

Morphometric Measurements

Morphometric measurements were made following Hubbs and Lagler (1970). Characters chosen included those previously used in the segregation of the three groups such as the length of 2nd and; 3rd dorsal spines, body depth, and head length plus numerous other characters. In all, 52 morphometric measurements (Table I) and 13 meristic counts
(Table II) were made on each fish. Standard length measurements were made to the nearest mm . All other measurements were made to the nearest 0.01 mm with electronic digital calipers. Each variable was measured four times and the average reading was recorded. Meristic variables were also counted four times and the mean recorded for each variable. Meristic counts such as number of scale rows above, below, and on the lateral line, gill raker counts, and fin ray counts were made under magnification.

Traditionally, data collected for morphometric analyses have been analyzed without much regard for allometry or the variation of allometry within populations. Growth and body form have been studied mostly by analysis of length or height and such measures have been criticized for being of limited utility in discerning shapes from sizes. Conventional measures of length commonly used in morphometric studies have been found (Bookstein et al. 1985) to be (i) aligned mostly along a few axes usually longitudunal or 'anterior-posterior' and with limited data on depth or breadth, (ii) restricted to some regions of the body with other regions receiving limited or no coverage, (iii) highly repetitious of some morphological landmarks such as the tip of the snout and end of vertebral column. Errors can also be made if measurements, (iv) are reliant on 'extremal' measurements which can be quite variable such as maximum body depth, and (v) cover too long a span thereby containing less information on localized variation than would relatively short distance measures. To overcome
these objections, morphometric variables chosen for analysis in this study (Table I) represented all regions of the fish including longitudunal, tranverse, and oblique axes.

Previous studies on the identification of Morone sp. have relied heavily on ratios derived from morphometric analyses for conclusive separation. For example, Williams (1976) determined that ratios of body depth/head length and fork length/body depth were conclusive in the separation of the striped bass $x$ white bass hybrid from either parental species. The use of ratios of morphometrics is one of three methods used to compare shapes among groups while removing the effects of size. The other methods are regression and principal component analyses. Although the use of ratios have been a long established technique for analyzing morphometric data, its reliability has been questioned because spurious variations can occur as a result of correlations between denominators and numerators as well as dependence of numerator on denominator (Atchley and Anderson 1978; Mosimann and James l979; Hills l978; Humphries et al. 1981). Consequently, use of ratios in morphometrics is being discouraged and methods which rely less on ratiorelated measures are being encouraged. One technique commonly used to nullify the effects of size in morphmetrics has been log-transformation of data. However, log transformation does not automatically remove size-related effects on variables (Humphries et al. 1981). To overcome these difficulties, it is now recommended that multivariate tech-
niques be used to make comparison.
Selection of variables to be included in a multivariate analysis is critical and has been based mostly on the criterion that a large number of variables be used. Considerably less importance has been attached to the functional and ecological roles of the variables chosen. I have attempted to include large numbers of variables but also to consider function and ecology in my selection of variables. In addition, $I$ have attempted to minimize redundancy in choice of variables and to include variables from all regions of the body (Table I, II). Differences in the morphometrics between the hybrid and the parental species, if they did exist, were expected to be small hence $I$ selected a large number of morphometric variables, encompassing most regions of the body. Many statistical tests assume homoscedasticity (equality of variance) and additivity of variance components. I do not believe this is a valid assumption, therefore, $I$ used variable transformations and tested the distributions of the $\log$ transformed variables for goodness of fit to the normal distribution prior to use in the model. All morphometric variables were logtransformed (unless otherwise indicated to base 10) before use in the analyses (Table III).

Statistical Analyses

The weaknesses inherent in univariate methods and ratios for taxonomic analyses caused me to use a multi-
variate approach employing discriminant analyses, principal component analysis, and sheared principal component analysis algorithms for separating the three groups. In recent years both classical and numerical taxonomists have employed discriminant analysis in the assessment of morphometric divergence among natural populations. Discriminant analysis employs a dependent variable (species type) and a collection of independent variables, including both continuous (morphometrics) and discrete (meristic) variables. It is recommended that separate analyses be performed on morphometric and meristic variables (Bookstein et al. 1985). In my study, there were high overlaps in the ranges of the meristic variables for all three groups. These high overlaps made statistical analyses impossible; consequently, further analyses of the meristic data were discontinued. The result presented here are based entirely on morphometric data.

The discriminant functions used in this study are developed in a multidimensional space that maximizes the variability between groups (in this case species) while minimizing the variance within groups. The discriminant analysis then develops a number of canonical variates (eigen vectors) which discriminate the groups. Each variate makes its own individual contribution to the discrimination, the first contributing the greatest amount, the second the next greatest, etc. In presenting the data, the means of each of the groups are plotted along the axes of the first two or three canonical variates in order to
depict the discrimination patterns. The discriminant function in its final form becomes a very powerful tool that uses the joint probabilities of combinations of variables for the best separation of the groups.

In my study, a multivariate analysis of variance (MANOVA) was first performed on the morphometric variables to test the null hypothesis of equality of the group centroids. Although MANOVA is a simple linear test, it should be conducted prior to multivariate discriminatory techniques to ensure that discrimination is not spurious. Discrimination is valid only if there is a significant difference among the group centroids (Pimental 1979). To accomplish this separation of my data, all 52 morphometric variables were analyzed iteratively using the step-wise linear discriminant analysis program BMDPO7M (Jennrich and Sampson 1983). The discriminant function is constructed in a step-wise manner such that at each step the relative contribution of each variable is assessed and if it falls below some criterion level, it is deleted from the analysis. Variables which make the largest contribution to the separation of all the groups (based on the $F$ value it generates in a one-way ANOVA) are entered first. After the first variable is entered into the discriminant function, the second variable making the next largest contribution (the one with the next largest adjusted $F$ value) is selected, etc., until all variables are used. Weights are then assigned to each variable such that the maximum difference between the groups is produced. Statistical proce-
dures employed in the discriminant analyses are described in detail by Green (1978), Morrison (1967), and Srivastava and Carter (1984).

In addition to indicating the relative contribution made by each variable to the overall discrimination, the scriminant function coefficient was also used to determine a variable's effect on group membership. When the value of a variable with a positive corresponding function is increased, the resultant discriminant score is increased. This tendency increases the probability of classifying the individual into the group with a higher centroid. On the other hand, increasing the value of a variable with a negative corresponding coefficient results in increased probability of classifying the individual into a group with a lower centroid.

I also used linear discriminant function analysis to test the reliability of my identification models. In applying the linear discriminant function analysis, the function is first developed, and second tested, based on the percent correct/incorrect classification of an independent sample of known specimens. The higher the percent of correct classification, the more useful the function is as a tool in the discrimination of the specimens. Individuals are assigned group membership based on the value of the classification function score. The classification function $C_{i}$ for species i is given by:

$$
c_{i}=c_{i 1} v_{1}+c_{i 2} v_{2}+c_{i 3} v_{3}+\ldots+c_{i j} v_{j}+c_{i p} v_{p}+c_{i 0}
$$

where the $C_{i j}$ 's are the classification function coeffi-
cients, $C_{i 0}$ is a constant, and the V's are the discriminating variables. An individual is classified into the group for which it's set of $V$ 's produces the highest classification score.

Discriminant analysis is an €scellent technique for depicting multivariate patterns of differences among the species but tells little of the interrelationships of the characters' overall groups. I used techniques such as factor analysis or principal component analysis (PCA) which generates factors which are often interpretable in terms of functional groups of variables to obtain such information. Factor scores may be generated for each individual, and these scores represent essentially new characters on which ANOVA and other tests may be done.

I also used principal component analysis (PCA), an ordination technique used for the analysis of the structure of multivariate observations, for investigating the morphological variations occurring in the populations. PCA has been widely used as an exploratory technique for discovering structure in morphometric and meristic data (Blackith and Reyment 1971; Smith 1973; Thorpe 1976; and Pimental 1979) and is particularly useful when no a priori patterns of interrelationships can be suggested or are suspected (Blackith and Reyment 1971). When used on morphometric and meristic data, PCA can be used to show geographic, clinal and other types of variations. The technique may be used on individual specimens since the analysis assumes very little by way of biological models (Thorpe 1976).

The goal in PCA is to reduce the original number of variables to a smaller number of variates or principal components which are expressed in terms of linear combinations of the original variables. The new variables (principal components or PCs) define independent patterns of variation among the original variables, which may be interpreted separately. The principal components are ordered in descending order of contribution to the explained variation such that $P C$ I accounts for the maximum percentage of variance possible for the single linear combination of the original variables. Principal component II is uncorrelated with principal component $I$ and represents a set of variables which explains the second largest amount of variation, and so on.

Shape is important in fish systematics and when used in conjunction with meristic charcters and color patterns, can serve as a very important tool for systematics and taxonomy. For example, white bass are described as being relatively more 'deep bodied' than striped bass of comparable length (Bayless 1972). I used principal component analysis to consider the effects of both size and shape since these two characters are often confounded (Humphries et al. 1981). In the analysis of a single population, the approach is quite simple because PC I is considered to be a general size factor. However, in the analysis of multigroup populations, size and shape effects often occur in the first two principal components (Humphries l984). Consequently, an interpretation of a multigroup analysis as we
have in my data would be confounded by size-dependent differences. In order to nullify the effects of size variability in morphometrics, it is necessary to develop a size-free PCA by partitioning the size effects from PC I and PC II. The result of the removal of size-related effects from PCA results in analysis of variation in shape. The technique for developing a size invariant PCA is called 'shearing' (Humphries et al. 1981). I used sheared PCA and discriminant analysis in the morphological separation of the three groups in this study.

The sheared principal component, $H$, is a linear combination whose coefficients equal the partial covariances adjusted for intergroup size. The value $H$ is approximately equal to the residual obtained from PC II after regressing out intragroup size, and is uncorrelated with intragroup size.

The need for shearing is determined by examining the plot of the regular principal component scores obtained from a covariance matrix for the presence of the same trend for PC I and PC II. Such a trend is an indication of sizerelated effects and necessitates corrective action through shearing. Shearing is accomplished by first standardizing the scores from the original PCA to zero mean, yielding two new components $P C I_{z}$ and $P C I_{z}$. The magnitude of the size effect is then computed by regressing PC IIz on $S$, the within group size component of $Q$ (the covariance matrix adjusted to mean zero within each group). The estimated residual from the above regression gives an estimate of $H$,
the sheared principal component. $H$ can then be used as a size invariant morphometric discriminator while still maintaining all of the original variance.

Size was an important confounding effect in my data, therefore a sheared principal component analysis was performed on the log transformed morphometric data using a SAS macro program Bookstein et al. (1985). The principal components were computed using the covariance matrix method.

Results

Discriminant Analysis of Striped bass,
White bass, and Striped bass $x$

## White bass hybrids

Results of the threegroup linear discriminant analysis, based on 52 morphometric variables, indicate that white bass, striped bass, and striped bass $x$ white bass hybrids are statistically separable. The seven variables which contributed significantly to the discrimination (Table IV) are:
(i) caudal peduncle length
(ii) snout length
(iii) first dorsal fin base length
(iv) pelvic fin base length
(v) pectoral to second dorsal fin distance
(vi) pectoral fin height
(vii) internasal distance

Application of stepwise discriminant analysis algorithm (BMDP07M) to unstandardized data results in a ranking of the variables in descending order of their contribution to overall discrimination. The analysis (Table IV) indicates that caudal peduncle length was the most important variable for separating the three taxonomic groups. Snout length was the next most important variable. A plot of the canonical variate or discriminant scores (Table V) shows the presence of three 'clusters' representing the three groups (Figure 1). Interpretation of the clusters in terms of the discriminant functions and the appropriate morphometric variables reveals that striped bass have longer snout lengths, caudal peduncle lengths, and internasal distance than do the other groups. Furthermore, the information indicates (Table IV; Figure l) that for each variable along the first canonical variate, the relative morphology of the hybrid is intermediate between those of both parental species. Fish with longer pectoral fin to second dorsal fin distances have higher probability of being classified as white bass whereas those with longer snout lengths are more likely to be classified as striped bass. The group centroids also indicate that complete segregation occurs along the first canonical axis but only partially along the second axis (Table VI). Canonical variate $I$ explains the most variation (77.71\%) and the variables which load high on this axis are the most important for group separation.

Although partial separation of the three taxonomic groups was achieved by the first discriminant function, the
second function only enables the separation of the striped bass $x$ white bass hybrid from both parental groups but not the separation of the parental groups from one another (Table IV; Figure 1). The first discriminant function accounts for $77.71 \%$ of the variation (variance $=A_{1} / A_{1}+A_{2}$ where $A_{1}$ and $A_{2}$ are the eigen values for functions $I$ and I respectively) while function $I I$, which separates the hybrid from the two parental forms, accounts for $22.29 \%$ of the variation. Function II indicates that specimens with long base length of first dorsal fin have a higher probability of being classified as hybrids (being hybrids) than as either of the parental groups. The canonical correlation $p$ of function $j$, (where $p_{1}=A_{1} / 1+A_{1}$, and $A_{l}=e i g e n$ value for function $I$ ) (the correlation between the linear discriminant function and the original set of variables) is 0.9837 for function $I$ and 0.9564 for function I (Table IV).

The results of the classification indicates that a 100\% correct classification into the pre-determined number of groups (reference samples of known identities) was obtained based on the seven morphometric variables (Table VII). Although discriminant function $I$ accounted for only $77.71 \%$ of the total variation, a complete separation of the three taxonomic groups was achieved by the model using the seven variables. Based on the results of this discriminant analysis (Table VIII) the following function was developed to classify any unknown fish into one of the three groups. Group I=

$$
\begin{aligned}
& \quad-300.36035 * \mathrm{~L}_{1}-2926.40112 * \mathrm{~L}_{2}-1236.32837 * \mathrm{~L}_{3}-453.61182 * \mathrm{~L}_{4} \\
& \quad+6003.64063 * \mathrm{~L}_{5}+3447.52490 * \mathrm{~L}_{6}-3738.73706 * \mathrm{~L}_{7}-3324.34546
\end{aligned}
$$

$$
\begin{aligned}
& -245.57773 * \mathrm{~L}_{1}-2666.80835 * \mathrm{~L}_{2}-737.22559 * \mathrm{~L}_{3}-558.96631 * \mathrm{~L}_{4} \\
& +5150.37891 * \mathrm{~L}_{5}+2942.44629 * \mathrm{~L}_{6}-3164.17896 * \mathrm{~L}_{7}-2654.37329
\end{aligned}
$$

Group III=

$$
\begin{aligned}
& 7.58869 * L_{1}-2422.22412 * L_{2}-1069.62012 * L_{3}-766.17554 * L_{4} \\
& +5120.94141 * L_{5}+3224.96753 * L_{6}-3350.47729 * L_{7}-2792.80615
\end{aligned}
$$

where $L_{1}-L_{7}$ respectively represent the log transformed values of the variables in Table IV. In this analysis, an individual is classified into one of the three species depending on the relative magnitude of the classification function, thus: If the Group $I$ coefficient is greater than Groups II and III coefficients, then the sample is classified as a white bass. If the Group II coefficient is greater than Groups I and III coefficients, then the sample is classified as a hybrid. If the Group III coefficient is greater than Groups I and II coefficients, then the sample is classified as a striped bass.

In order to test the accuracy of the model, the classification function was applied to an independent sample "holdout sample" of Morone sp. from several oklahoma reservoirs. The result (Table IX) indicates that a $91.67 \%$ discrimination was obtained between the three forms. The classification scores indicate that the white bass and striped bass $x$ white bass hybrids are morphologically very similar (Table $X$ ). The result also confirms the presence of two or more Morone sp. in some Oklahoma reservoirs.

Sheared Principal Component Analysis of White bass, striped bass, and striped bass $x$ White bass hybrids

The results of principal component analysis performed on the log transformed morphomertric variables indicated the presence of three 'clusters', representing the three taxonomic groups (Figure 2). However, the scatter plot of $P C I$ against $P C$ II showed the presence of size-dependent distributions. Plots of $P C I I$ and $P C$ against log of standard length also indicated that $P C I$ (Figure 3) and PC II (Figure 4) were both size confounded.

In order to achieve a size-free discrimination of the three groups, I sheared, (Humphries et al. 1981) the data to obtain a sheared principal component, $H$, (Table XI) which when plotted against $P C I$, removes the effect of size. After this treatment, the data on striped bass had significantly different shape components than white bass and striped bass $x$ white bass hybrids but the latter two did not differ from one another (Figure 5). The strong morphological similarity between the white bass and the hybrid was indicated by the mean sheared PC II which are -0.358, 0.396 , and -0.704 for the white bass, hybrid, and striped bass respectively. However, these values also indicated that the hybrid was morphologically intermediate between the parental species.

Morphometric similarity between white bass and hybrid
was also indicated when the sheared PC II was compared with the results of the discriminant analysis. Canonical discriminant analysis indicated (Figure l) that the three groups could be separated using variables associated with canonical variate I. However, a plot of sheared PC II plotted against the canonical variate $I$ indicated (Figure 6) that, although white bass and hybrid can be distinguished morphologically from the striped bass, they can not be distinguished from one another. The correspondence between the results of the discriminant analysis and sheared principal components is strong (Pearson correlation coefficient $=0.89$, $p<0.0001$ ) suggesting that the results of the discriminant analysis were not significantly confounded by size. A similar examination of the relationship between sheared PC II and canonical variate II of the discriminant analysis also indicated (Figure 7) that the white bass and hybrids are statistically similar morphologically but they are collectively different from the striped bass (Pearson correlation coefficient $=-0.36, \mathrm{p}<0.04)$. The above comparisons between both multivariate classification techniques suggest that subtle differences do exist in the morphology of the three groups, but that white bass and hybrids are morphologically very similar. These analyses also suggest that differences between them may not be discernable when examined in conjunction with the striped bass.

Sheared Principal Component Analysis<br>for White bass and Striped bass $x$<br>White bass hybrids

Computation of principal components on white bass and hybrids indicates that although an oblique 'cluster' does exist due to some size confounding on PC I, it is much less pronounced in the absence of striped bass (Table XII, Figure 8) than it was with striped bass included in the data set. Shearing of $P C$ II of this reduced data set removed the effects of size on the data and resulted in a complete morphological separation of the white bass and striped bass $x$ white bass hybrids. This elimination of size as a factor is evidenced by the size invariant pattern exhibited by the plot of sheared PC II against log of standard length (Figure 9). A plot of the sheared PC II against PC I (Figure 10) also illustrates the ability of this procedure to separate the two groups.

In summary, white bass can be separated from hybrids on morphological grounds if striped bass are not included in the analysis. The inclusion of striped bass in the data set results in differences between white bass and hybrids being masked.

Discussion

The introduction of striped bass $x$ white bass hybrids
into reservoirs with native populations of striped bass and (or) white bass has created successful sport fisheries in several Oklahoma reservoirs. However, these introductions have also generated controversies. Some individuals believe that hybrid introductions have adversely inpacted largemouth bass and white bass populations (K. E. Erickson, ODWC). Conversely, there is some evidence (Gilliland and Clady 1981) that only limited impacts have occurred.

The introductions of hybrids into warm water reservoirs have often been justified on the basis of the hybrid's ability to utilize the clupeid forage without considering the effects on other game species. However, in reality, a thorough understanding of the biological interactions between these groups is necessary before realistic management decisions can be made. The first requirement for understanding the interaction between these groups is the ability to differentiate hybrid stocks from those of the parental groups. In addition, the ability to distinguish the hybrid from the white bass, the parental group to which it is morphologically more similar, has obvious management implications. For example, in Oklahoma there are no creel limits on white bass and striped bass $x$ white bass hybrids in most reservoirs. Thus it is imperative that the angling public as well as the fisheries biologist and game ranger be able to distinguish the two forms from the striped bass for which there is a creel limit on most Oklahoma reservoirs. The results of this study indicate that this separation is not easy and that with multivariate examination of

52 morphometric variables, the precision of the separation of the hybrid from the white bass is low. Inclusion of striped bass in the analysis further complicated the separation.

In spite of the difficulties of the separation of these forms, there are significant differences in the morphometric measurements for white bass, striped bass, and striped bass $x$ white bass hybrids. Separation of the forms is possible with sheared principal component analysis and discriminant analysis. Estimates of the mean sheared PC II between the three groups indicate that striped bass is most different from the white bass and also that high similarity between the hybrid and white bass results in less reliable predictions of group memberships when all three groups are present. The similarity between these two groups as indicated by the mean sheared PC II between the hybrid and white bass is confounded by size-related factors but the high correlation ( 0.89 ) obtained between the size-free PCA and the discriminant analysis indicates that the results of the discrimnant analysis (unadjusted for size) are reliable and that the effects of size on the overall morphological discrimination was probably minimal.

Other authors using univariate approaches have found the hybrid to be more similar in some characteristics to either one or the other of the parents. For example, williams (1976) determined that the mean ratio of body depth/head length of the hybrid was closer to that of the white bass than to that of the striped bass and also that
the mean fork length/body length of the hybrid was also closer to that of the white bass than to that of the striped bass. Conversely, he found that mean head length/2nd anal spine length was more similar between the hybrid and striped bass than between the hybrid and white bass.

The multivariate analysis also shows that the hybrid is morphologically intermediate in some characteristics between both parental forms, but also allows us to determine what such differences mean. The distance between the pectoral fin and the second dorsal fin was longer on the average for white bass than for the other two groups and the hybrid was intermediate. This observation means that measures of shape along oblique axes are important variables for distinguishing the three groups. The longer pectoral fin to second dorsal fin distance for the white bass coupled with its greater tendency to acquire longer pectoral fin height and longer base length of pelvic fin are all indicative of morphological developments associated with the humeral region of the white bass.

That the morphological differences separating the white bass from the other groups is the greater development in the humeral region, is also corroborated in part by univariate determinations. Williams (1976) concluded that the mean body depth/head length ratio for white bass and hybrid was 1.198 and 0.893 respectively. These findings indicate that white bass may have a more developed humeral region than the other two groups. This greater development
is reflected in the greater body depth for the white bass. The joint effect of the variables around the humeral region could be construed to indicate that there is a difference in the degree of rotundness of the abdominal and humeral regions of the white bass relative to those of the other two groups. This conclusion is supported by discriminant analysis because the intermediate value of the loadings for the hybrid along canonical variate II are also a relative measure of body depth and rotundness. Although canonical variate scores for the white bass indicate differences in growth and body morphology associated with rotundness and increased condition factor (a measure of allometry) relative to the striped bass, the data also indicates that striped bass have longer caudal peduncle length, snout length, and internasal distance than the white bass.

The association of variables such as caudal peduncle length and snout length on one axis imply that striped bass are more streamlined than the other two groups. Some predators such as striped bass which actively chase down their prey are characterized as having 'torpedo-shaped' fusiform body form and well developed caudal region for rapid propulsion (Moyle and Cech 1982). Conversely, the white bass which is presently found in reservoirs through out the midwest and in most of the United States (Trautman 1981) is primarily a stream dwelling fish which has been widely introduced or entrapped in reservoirs. In keeping with these differences in behavior, white bass are less fusiform and more rotund. One might hypothesize that a com-
parison of the predatory behavior of the white bass would reveal investment of considerably different amounts of time, energy, and effort in foraging.

Conclusions and Recommendations

The popularity of the striped bass fishery in the Chesapeake Bay, on the east coast of the United States, and in landlocked reservoirs, coupled with the importance of the white bass fishery, resulted in the development of the female striped bass $x$ male white bass hybrid. Introduction of the hybrid has been encouraged because of its ability to utilize the clupeid forage abundant in several reservoirs, its 'fighting' ability, its rapid growth, and ability to adapt to different geographic regions and water quality conditions.

Although the hybrid fishery is over 20 years old, questions are still being raised about the ecology and management of the hybrid. The most important questions raised by the introductions into numerous streams/reservoir systems are (i) the potential ecological impacts on native populations of white bass, striped bass, largemouth bass and other species, and (ii) management problems stemming from uncertainty in stock/species identification in aquatic systems in which the hybrid coexists with one or both of the parental species. Since sound fisheries management entails accurate stock identification and assessment, the importance of unambiguous and accurate techniques for stock
separation are a nesessity.
Separation of the three groups have traditionally been done on the basis of size since striped bass is the largest and white bass the smallest. Identification was also done using meristic and morphometric characters such as the number and shape of tooth patches on the base of the tongue as well as the shape and patterns of stripes on the sides of the fish and measurement ratios. In Oklahoma, these methods have been proven unreliable for separating the three groups, especially when small fish are involved. Thus a multivariate approach was employed as an aid in identification.

Separation of the three groups was possible but difficult when all three groups were analyzed simultaneously with discriminant analysis and sheared principal component analysis. When data on white bass and hybrid were analyzed the absence of data on striped bass, separation was easier. The variables which contributed significantly to the discrimination were the caudal peduncle length and the snout length. The base lengths of the first dorsal and pelvic fins were longer on the average for the hybrid and ratios involving these measurements can be used to separate the hybrid from both parental species. White bass tended to have longer pectoral to first dorsal fin distances than the other two groups whereas striped bass tended to have longer caudal peduncles and snout lengths than the other two groups. For the hybrid, the range of the above variables tended to be intermediate between those of the parents.

These variables can be used to separate the three groups when they coexist.

The application of the discriminant analysis to an independent data set, comprised of fish from several Oklahoma reservoirs, confirmed the presence of stocks of two or more Morone sp. in some reservoirs. Fort Supply reservoir, Canton reservoir, and Lake Carl Blackwell had stocks of both white bass and striped bass $x$ white bass hybrids whereas Optima reservoir contained only striped bass $x$ white bass hybrids.

Management implications of the findings are that fishing regulations, regarding creel limits on the three Morone forms, need to be reevaluated especially in lakes which contain the striped bass and one or both of the other forms. This is necessitated by the lack of a creel limit on white bass and striped bass $x$ white bass hybrids in most reservoirs whereas there is a creel limit on striped bass in most reservoirs. The strong similarity between the three forms and their occurrence in the same bodies of water could result in high incidences of misidentification which would complicate management objectives. The inability to accurately and easily identify the white bass from the hybrids in Oklahoma waters makes enforcement of game laws difficult. Such a problem can be alleviated by managing the two species as a single fishery. Benefits from management of a mixed Morone sp. as a single fishery rather than as separate fisheries are probably high. However, application of the classification function developed in this study
(91.67\% separation) suggests that the risks associated with misclassification may be acceptable in some situations. In those situations, the single-species management concept should be used. The need for joint management of the white bass and the striped bass $x$ white bass hybrids as one fishery was recognized by the ODWC whose management guidelines (1987 Oklahoma fishing regulations) apparently addressed the problem by setting equal creel limits on both species.

Stocking the hybrid in reservoirs containing the parental species should be carefully considered because of difficulties in management of the fish populations and enforcement of game laws. Stocking of hybrids in reservoirs containing populations of white bass would entail managing both species as a single fishery (as suggested by the 1987 Oklahoma fishing regulations). Without easy methods for delineation of the various stocks and populations, management and law enforcement become problematic. Multivariate techniques are required for accurate identification of the three groups but these techniques are too complex to be used by the angling public or law enforcement officers. For these groups, the seven most distinguishing factors identified in the multivariate analysis should be used to assign group memberships. Ultimate group membership, however, can only be decided based on the value of the linear classification function coefficients.

## CHAPTER III

SIMULATION MODEL OF THE POPULATION DYNAMICS OF WHITE BASS AND STRIPED BASS X WHITE BASS HYBRID IN LAKE<br>CARL BLACKWELL, OKLAHOMA

## Literature Review

Historically, interest in population dynamics models has centered on generalized, theoretical population systems. In recent years however, there has been considerable interest in modeling specific fish populations, with the practical objectives of understanding or predicting population changes (Orth 1977, Taylor 1981), or simulating the behavior of populations under various management regimes (Reed 1982, Gutreuter 1983, and Zuboy and Lackey l975). Diverse approaches have been applied to these ends ranging from strictly empirical equations geared toward predictions (Orth 1977, Taylor 1981), to highly detailed biologically explicit models oriented towards understanding and simulation (Gutreuter 1983; Jester et al. 1977). Empirical models can be quickly constructed from raw data with little biological know-how, may have good predictive powers under conditions similar to those in which
the data were collected, but they add little to our understanding of biological processes. Highly detailed models require considerable biological knowledge and intuition, as well as large quantities of data, kut their complexity and extensive mathematical structure make interpretation difficult. Grant (1986), in his critique of large-scale systems models, proposes that the most important attribute of a model is transparency, or ability to provide insight into the workings of the system with a reasonable expenditure of effort. To maintain this attribute, Grant suggests that models be aimed at specific objectives, be simple, and be constructed with a balanced blend of fact, theory, and intuition.

Natural resource conservation agencies in several southern states, including the Oklahoma Department of Wildife Conservation, stock hybrids into reservoirs with native populations of white and (or) striped bass. The original purposes of these stockings were to (i) provide an alternative pelagic fishery to the white bass and (or) striped bass, and (ii) utilize the abundant clupeid forage present in most of these reservoirs to provide a harvestable resource. Most hybrid stockings in Oklahoma reservoirs are done every 2 to 4 years (Hicks 1978, l979; Kleinholz 1985; Maughan et al. 1985). Creel surveys and fall gill net sampling are used to assess the standing crops and other pertinent statistics of the hybrid and white bass populations.

To provide an alternative for predicting the charac-
teristics of these populations, I have developed a conceptual model of the population dynamics of striped bass $x$ white bass hybrids and white bass in reservoirs. I have patterned my efforts after Orth (1977), Reed (1982), and Taylor (1981). Orth (1977) and Reed (1982) developed models for population dynamics of largemouth bass in reservoirs and Taylor (1981) modeled walleye and trout population dynamics. Although these models represent "the state of the art" in modeling single-species reservoir fisheries, they are not applicable to multispecies fisheries such as the white bass and striped bass $x$ white bass hybrids. The goal of this model is to develop a tool for fishery biologists engaged in the management of striped bass $x$ white bass hybrids and white bass.

The major assumptions of this model were that densitydependent effects on growth and recruitment as well as random environmental fluctuations were in effect. Consequently, population sizes of both species were constrained around pre-determined densities. Gulland (1983) criticized single species models for neglecting the interactions between species. Because white bass and hybrids are ecologically similar, often occur in the same reservoirs, and are sometimes managed jointly, I decided to model both populations under the assumption that density-dependent mechanisms (especially forage) limit the populations. Environmental fluctuations are important in regulating fish populations. In reservoirs, these fluctuations often occur in the form of variations in water level and has been shown
(Zweiacker 1972; Orth 1977) to regulate largemouth bass recruitment in Lake Carl Blackwell. I assumed that white bass and hybrid populations in Lake Carl Blackwell were subject to random environmental fluctuations, hence $I$ introduced stochastic perturbations on the recruitment processes.

A good model is one which maximizes any two of the following properties: realism, precision, and generality (Levins 1968). Therefore, consistent with the major objectives of this study, many of the data used in implementing the model and the conclusions drawn there from are specific to striped bass $x$ white bass hybrids and white bass in Lake Carl Blackwell, Oklahoma. However with appropriate data, the model $c a n$ be adapted to simulate the population dynamics of a variety of two species fisheries consisting of a put-and-take fishery (such as the hybrid) and a naturally reproducing species.

The Model

The model $I$ developed utilizes information from several existing models including those of Jacobsen (1983), Orth (1977), Zagar and Orth (1986), Taylor (1981), Reed (1982), Gutreuter (1983), and Walters (1969). The model is age-structured, and stochastic and utilizes the Leslie matrix algorithm (Leslie l945) to express year class strengths.

Recruitment is generally defined as the number of fish
entering the harvestable size class during the given year and has been shown (Walters 1969 , orth 1977) to be the driving factor in fish population dynamics. Dynamic pool models have shown a very strong relationship between the size of the parental stock and the number of recruits. A number of factors affect recruitment including spawning biomass, environmental factors, predation, and competition (Pitcher and Hart 1981). In addition, Nikolskey (1969) stressed the importance of population size; indicating that changes in the number of recruits are responses to factors such as fecundity, viability of eggs and larvae, mortality, and growth all of which in turn vary in a complicated manner in response to environmental changes. These factors can be grouped as either density-dependent or densityindependent (Lackey and Hubert 1979).

Data on density independent factors such as environmental variability were not available for use in this model thus a density-independent stochastic component was used to estimate these factors. Long-term population statistics were also not readily available for white bass and striped bass $x$ white bass hybrids. As a result of paucity of data on these species, the resulting model is mainly a conceptual model with the theoretical framework developed for a multispecies fishery. The model is intended to serve as a guide to fishery biologists and policy makers regarding management of these resources.

A fish population will change in biomass after one time period as a result of recruitment, growth in biomass,
and mortality. This relationship can be expressed as

$$
\begin{equation*}
\mathrm{dB} / \mathrm{Bdt}=\mathrm{R}(\mathrm{~B})+\mathrm{G}(\mathrm{~B})-\mathrm{M}(\mathrm{~B})-\mathrm{F}(\mathrm{E})+\mathrm{e} \tag{1}
\end{equation*}
$$

where $B=b i o m a s s, E=f i s h i n g$ effort, $R, G, M$, and $F$ are rates of recruitment, growth, natural mortality, and fishing mortality and $e$ is a random error. Derivations of equations 2 through 25 follow the methods of Orth (1977), Ricker (1975), Pitcher and Hart (1981), and Gulland (1983). The instantaneous rate of total mortality (Gulland 1973) can be written as

$$
\begin{equation*}
d N_{i j} / d t=-Z_{i j} N_{i j} \tag{2}
\end{equation*}
$$

 class $j$ of species $i$, and $N_{i j}=$ population size of age class j of species i. Rearranging equation (2), we can obtain

$$
\begin{equation*}
d N_{i j} / N_{i j}=-Z_{i j} d t \tag{3}
\end{equation*}
$$

Integrating (3) yields

$$
\begin{equation*}
N_{i(j+1)}=N_{i j} \exp -(Z i j) \tag{4}
\end{equation*}
$$

$Z_{i j}$ can be partitioned into fishing mortality, $F_{i j}$, and natural mortality, $M_{i j}$, thus

$$
\begin{equation*}
N_{i(j+1)}=N_{i j} \exp ^{-(F i j+M i j)} \tag{5}
\end{equation*}
$$

Instantaneous rate of survival can be obtained from the following relationship

$$
\begin{equation*}
S_{i j}=N_{i(j+1)} / N_{i j} \tag{6}
\end{equation*}
$$

alternatively, $S_{i j}$ can be derived from instantaneous total mortality rate as

$$
\begin{equation*}
S_{i j}=\exp ^{-(Z i j)} \tag{7}
\end{equation*}
$$

where $S_{i j}=i n s t a n t a n e o u s ~ a n n u a l ~ s u r v i v a l ~ r a t e ~ f o r ~ a g e ~ c l a s s ~$ $j$ of species i. Instantaneous annual rate of growth, $G_{i j}$, was computed from the formula

$$
\begin{equation*}
G_{i j}=L_{i(j+1)} / L_{i j} \tag{8}
\end{equation*}
$$

where $L_{i(j+1)}=$ mean total length of age class $j+1$ of species i, and $L_{i j}=m e a n$ total length of age class $j$ of species i. Growth in weight can be expressed as

$$
\begin{equation*}
W_{i(j+1)}=W_{i j} * \exp (b i * G i j) \tag{9}
\end{equation*}
$$

where $W_{i}(j+1)=$ mean weight of age class $j+1$ of species $i$, $W_{i j}=$ mean weight of age class $j$ of species $i$ and $b_{i}$ is the exponent in the length-weight relationship

$$
\begin{equation*}
\text { Weight }=a_{i} * \text { Length }{ }^{b i} \tag{10}
\end{equation*}
$$

Biomass $B_{i j}$, of age class $j$ of species $i$ was computed from the formula

$$
\begin{equation*}
B_{i j}=N_{i j} * W_{i j} \tag{11}
\end{equation*}
$$

The mean number of fish in age class $j$ of species $i$ during the period $t$, $t+d t$ is represented by the formula

$$
\begin{equation*}
\operatorname{NBAR}_{i j}=\int_{t}^{t+1} N_{i j}^{*} \exp ^{-(z i j)} d t \tag{12}
\end{equation*}
$$

Integrating (12) yields

$$
\begin{equation*}
\operatorname{NBAR}_{i j}=(1-\exp (-Z i j)) / z_{i j} \tag{13}
\end{equation*}
$$

Mean biomass (kg) of age class $j$ of species $i$ during time period $t, t+d t$ is represented by the formula

$$
\begin{equation*}
\operatorname{BBAR}_{i j}=\iint_{i j}^{t+1}(\exp (G i j-Z i j)) * t d t \tag{14}
\end{equation*}
$$

integrating (14) ${ }^{\mathrm{t}}$ yields
BBAR $_{i j}=B_{i j}\left(1-\exp ^{-(Z i j-b i * G i j)}\right) /\left(z_{i j}-b_{i} * G_{i j}\right)$
when $Z_{i j}>G_{i j}$ and

$$
\begin{equation*}
=B_{i j}{ }^{*}(\exp (G i j-z i j)-1) /\left(G_{i j}-Z_{i j}\right) \tag{16}
\end{equation*}
$$

when $Z_{i j}<G_{i j} . B_{i j}$ is the biomass of age class $j$ of species $i$ at time $t=0$. Number of fish of age group $j$ of species $i$ harvested during the time period $t, t+1$ is represented by the formula

$$
\begin{equation*}
C_{i j}=F_{i j} * \operatorname{NBAR}_{i j} \tag{17}
\end{equation*}
$$

where $F_{i j}$ is the fishing mortality rate of age-class $j$ of species i. Weight of fish (kg) of age group $j$ of species i harvested during the period $t, t+1$ is represented by the formula

$$
\begin{equation*}
Y_{i j}=F_{i j} * \operatorname{BBAR}_{i j} \tag{18}
\end{equation*}
$$

Gross production of fish of age class $j$ of species i during the period $t, t+1$ is represented by the formula

$$
\begin{equation*}
\operatorname{GPROD}_{i j}=b_{i} * G_{i j} * \operatorname{BBAR}_{i j} \tag{19}
\end{equation*}
$$

Net production (Ricker 1975) by age class $j$ of species i during the period $t, t+1$ is given by

$$
\begin{equation*}
\operatorname{NPROD}_{i j}=\left(b_{i} * G_{i j}-Z_{i j}\right) * \text { BBAR }_{i j} \tag{20}
\end{equation*}
$$

Excess of growth over natural mortality of age class $j$ of species $i$ during the period $t, t+1$ is given by

$$
\begin{equation*}
\operatorname{ExS}_{i j}=\left(G_{i j}-M_{i j}\right) * \text { BBAR }_{i j} \tag{21}
\end{equation*}
$$

and weight of fish of age class $j$ of species $i$ that die naturally is represented by the formula

$$
\begin{equation*}
\mathrm{WTMORT}_{i j}=\mathrm{M}_{i j}{ }^{*} \text { BBAR }_{i j} \tag{22}
\end{equation*}
$$

There is a concensus of opinion that no general or specific parental-recruitment relationship holds true for all species (Beverton and Holt 1957, Ricker 1958). Furthermore, in situations where a theoretical basis for a particular model exists, it still shows significant dispersions; indicating that factors other than parental stock may be in operation (Reynolds and Babb 1978). This indicates that other factors such as stochastic environmental perturbations may be important in determining recruitment. Assuming that the above assertion is correct, I included a stochas-
tic component into the recruitment submodel. Biomass and other statistics of the fishery computed by the model include fish of both species within age classes $I$ through $V$. Data on hybrids older than age $V$, and larval and young-of-the-year fish were not available hence the model is also limited to age-classes I through V.

## Parameter Estimation

## Growth Rates

Growth in length is a function of many factors, the most important of which are initial size and prey availability. Within any given population, smaller fish grow at a much faster rate than larger fish and at any given size this rate is usually constant (Weatherly and Rogers 1978). Conversely, prey availability is unpredictable and accurate estimates of prey availability have been considered to be one of the weak points of single-species modeling (Reed l982). In any balanced fish community (Swingle 1950), predator numbers exert a controlling influence over prey numbers hence predators are limited by their own biomass. prey availability is a function of predator size and numbers, prey size and numbers, prey distribution, time, energy requirements, competition, and other factors (werner 1974). Since data on prey availability were not available for use in this model, density-dependent and stochastic effects of this factor on
growth, as estimated from fish in Lake Carl Blackwell, were used to build the model.

Mean lengths-at-age data were taken from data from fish collected from Lake Carl Blackwell during 1985 and were fit to von Bertalanffy growth curves (von Bertalanffy 1938) for each species. Walford plots (Walford 1946) were obtained by plotting mean lengths at time $t+1$ against mean lengths at time $t$. The slopes of the Walford plots equals exp(-ki) where $k_{i}$ is the asymptotic rate of growth for species i. Ricker (1975) and Cailliet et al. (1986) give detailed descriptions of the von Bertalanffy curve and how to estimate the parameters. The simplest version of the von Bertalanffy growth formula has the form

$$
\begin{equation*}
L_{i t}=L_{m i}\left(1-\exp ^{-k i(t i-t 0)}\right) \tag{23}
\end{equation*}
$$

$k_{i}$ is the asymptotic growth constant for species $i$ and $t_{0}$ is the 'age' the fish would have had at zero length if they had always grown according to the equation $\left(t_{0}\right.$ genearlly has a negative value). The maximum lengths attainable by each species, $L_{m i}$, were obtained from the Walford plots as the points of intersection of $L_{i}(t+1)=L_{i(t+1)}$ with $L_{i(t+1)}=L_{i t}$. The intercept, $I_{i}$, of the plot of $\log \left(L_{m i}-L_{i t}\right)$ against $t$ enables the computation of $t_{0}$, thus

$$
\begin{equation*}
t_{0}=\left(I_{i}-\log \left(L_{m i}\right)\right) / k_{i} \tag{24}
\end{equation*}
$$

The von Bertalanffy growth model for the white bass was

$$
\begin{equation*}
L_{i t}=448 *\left(1-\exp ^{-0.48(t i-0.13)}\right) \tag{25}
\end{equation*}
$$

and for the hybrid, was

$$
\begin{equation*}
L_{i t}=487 *\left(1-\exp ^{-0.53(t i-0.08)}\right) \tag{26}
\end{equation*}
$$

Mean lengths-at-age, and estimated annual instantaneous
rates of growth for white bass and striped bass $x$ white bass hybrid are listed in Table XIII.

## Length-Weight Relationship

Length-weight relationships were determined for both species by regressing the common logarithm of weight (grams) against the log of total length (mm). The estimated constants and $b$, obtained from the relationship weight=a*length ${ }^{\text {b }}$ are listed in Table XIV.

## Mortality Rates

Mortality in exploited fish populations comes from two sources: fishing and natural causes. In unexploited populations, natural causes are the sole source of mortality whereas in heavily exploited populations, fishing may account for a major part of the total mortality. Fishing mortality is a function of the catchability coefficient, $q$, and the fishing effort, f. Natural mortality estimates are less precise than fishing mortality because they can not be estimated independent of fishing mortality; thus the errors associated with fishing mortality estimates are carried over into natural mortality estimates.

Instantaneous total mortality rates, $Z_{i j}$, were estimated for both species by means of catch curves (Ricker 1977, Bagenal 1978) on 1985 data from Lake Carl Blackwell. The natural $\log$ of catch per unit of effort plotted against
age, yields a graph with both ascending and descending arms. The descending arm represents the fully recruited age classes and its slope at various ages are taken to represent the age-specific instantaneous total mortality rates. Data on annual instantaneous rates of fishing and natural mortalities were not available for either species. Consequently, total instantaneous mortality rates were partitioned on the assumption that the fishing mortality rates for both species in Lake Carl Blackwell were slightly lower than that for the hybrid in Sooner Lake (Glass 1982). Mortality rates employed in the model are listed in Table XV.

## Survival Rates

The survival rates for both species were obtained from the data used to compute instantaneous total mortality rates. Age-specific survival rate, $S_{i j}$ is given by

$$
\begin{equation*}
S_{i j}=\exp ^{-(Z i j)} \tag{7}
\end{equation*}
$$

The age specific survival rates computed for both species and used in the modeling are listed in Table XVI.

## Fecundity

Fecundity estimates for white bass were obtained from Baglin (1972), who determined age-specific fecundity rates for white bass in Lake Texoma, Oklahoma-Texas. Baglin's estimate of fecundity, derived from fish in the Washita river arm of Lake Texoma, was:

White bass in Lake Texoma (Baglin 1972) and Lake Carl Blackwell did not reach sexual maturity until age II. The age-specific fecundities shown in Table XVII were used in the model.

The Simulation

The model assumes an annual stocking of a fixed number of hybrid fingerlings and a self sustaining white bass population. The number of hybrid fingerlings stocked annually is at the discretion of the modeler but the survival of the stocked fish is controlled by stochastic variation of environmental parameters as well as density-dependent factors. Conversely, the white bass population is controlled by the survival of the fingerlings produced by natural reproduction of a population with a l:l sex ratio. Furthermore, in the modeling effort, I assumed that interactions with fish sf きcies other than between white bass and hybrids were negligible or inconsequential and that both fish species were initially absent from the reservoir. Thus 3,000 white bass fry and 10,000 striped bass $x$ white bass fingerlings were used as 'seeds' to initiate the simulation. The population dynamics of both species were simulated for fifty years and statistics such as abundance, catch, yield, and lengths-at-age were generated on an annual basis.

Since the value of any density-dependent function is
determined only when the value of the density is specified, it was necessary to determine the absolute size of the population (numbers, biomass) at equilibrium. To obtain these estimates of absolute population size at equilibrium, six stations in Lake Carl Blackwell were sampled monthly for 14 months (June 1984 to August 1985 excluding January). Sampling was done with gill nets, trap nets, and barrel nets. Constant monthly fishing efforts were expended using 12, 61 m nylon experimental gill nets (mesh sizes 2.54, 5.08, 7.62 , and 10.16 cm respectively). These efforts resulted in harvests of both species that were considerably less than those from other Oklahoma reservoirs (Table XVIII). These data indicate that the absolute population size of both of these species in Lake Carl Blackwell is quite small.

To accurately determine the total standing crop or biomass in a reservoir the size of Lake Carl Blackwell (area=1369 ha) would require extensive mark-recapture data or cove rotenone data and determination of catchability coefficients for each sampling technique. These data were not available, therefore standing crop was estimated based on the sampling data available. A Morone sp. standing crop estimate of 700 kg (for both species) was employed as the upper limit beyond which density-dependent forces become operative.

## Fry and Egg Survival

Since young-of-the-year survival is probably the key element determining population fluctuations in reservoir fisheries, and because weather often influences their survival, several modeling
attempts have been made to include stochastic elements in predicting young-of-the-year survival (Jensen 1975; DeAngelis et al. 1980). One problem of these attempts has been the inclusion of stochastic events in predicting the survival of a pre-reproductive age class which may alter the intrinsic rate of growth of the population. Goodyear and Christensen (1984) attempted to adjust for this bias by determining a correction factor for the bias in population growth introduced by the distribution of the random variation. Even with these adjustments, estimates of survival from egg to age $I$ is difficult to obtain. Vaughan and Saila (1976) used a Leslie matrix algorithm to compute survival rates from egg to age $I$ as:

$$
\begin{equation*}
S_{0}=1 / \sum_{i=1}^{k-1}\left(m_{i+1}\left({\underset{j=1}{i}}_{i=S_{j}}^{i}\right)\right. \tag{28}
\end{equation*}
$$

where $S_{j}=a \operatorname{specific}$ survival rate, $k=n u m b e r$ of age classes, and $m_{i}=a g e$ specific fecundity. In my study, the annual stocking rate for for hybrids was substituted for age-specific fecundity in equation (27) above. Random variation was incoporated into the term representing the probability of survival from egg through age 0 by the equation

$$
\begin{equation*}
S_{0}(t)=S_{0} * \exp (R(t)) \tag{29}
\end{equation*}
$$

where $S_{0}(t)=$ probability that eggs deposited at the beginning of the year will survive to age $I$, and $R(t)=r a n d o m$ number for year $t$. Initially a correction factor, C, determined by the method of Goodyear and Christensen (1984) was used to obtain a corrected survival rate of:

$$
\begin{equation*}
S_{0}(t)=S_{0} * \exp (R(t)+C) \tag{30}
\end{equation*}
$$

However, since the model was subject to density-dependent constraints the true effects of the correction factor were masked. Consequently, the correction factor, $C$, was dropped from the model. In its place, a subroutine developed by Orth (1977) was adapted for use in computing survival rates of age 0 white bass and hybrids.

## Stochasticity

The ability of any model to provide reliable predictions depends on several uncertaintities associated with the modeling process. Uncertaintities in modeling can result from three sources (i) errors in the model construction, (ii) uncertainty in the model parameters, and (iii) variability of the natural system (O'Neill et al. 1979). In this context the density-dependent models contain virtually no bias compared to those inherent in the densityindependent Leslie matrix models where wide deviations occur.

Understanding large-scale fluctuations due to variations in year-class strengths have been the recipient of
considerable effort (Ward and Larkin 1964). In some cases, year-class size has been determined by density-dependent processes within the population. However, random fluctuations and their effects on recruitment are also important. Adjustments for these random effects is of ten done by multiplying the survival factor by a random variable (Allan and Basasibwaki 1974).

In order to account for random variability in recruitment, a stochastic component was introduced into the model developed in this study. The survival rates of white bass eggs and stocked hybrid fingerling to age $I$ were then estimated (equation (29)) by multiplying the original rates. by the exponent of a normally distributed random variate of mean of 1 and standard deviation supplied by the user. Thus the modeler could incorporate some realistic degree of fluctuation in the young-of-the-year population sizes.

Density-Dependent Effects on Young-of-the-Year Survival

Most fish populations adjust to increased mortality or excessive survival through one or more compensatory mechanisms. Density-dependent mechanisms have been incorporated into Leslie matrix models for mortality rates (Walters 1969; LMS 1975) and stock recruitment (Ricker type) models (Hess et al. l975). In addition a model designed to predict the long-term impact of man-induced mortality on the population must incoporate some form of
negative feedback mechanism (Gulland 1983).
Three problems confront the modeler when the Leslie matrix is used to model fish populations. They are (i) selection of age-specific parameter(s) that are assumed to be density-dependent, (ii) selection of a function relating a density-dependent parameter to some index of population size, and (iii) selection of numerical values for the parameters in this function. Density-dependent effects can be modeled in terms of (i) recruitment, (ii) mortality (fishing and natural), and (iii) survival of age-class 0 . Survival rates for age class 0 are very difficult to obtain directly (Vaughan and Saila 1976;). However, young-of-theyear survival is the most important parameter in determining reservoirs fisheries population dynamics (Orth l977), and it is important to consider the effects of densitydependence on this factor.

In my model, the function which describes the densitydependent effects on white bass egg and stocked hybrid fry survival is given by:

$$
\begin{equation*}
F(S(t))=F(B(t)) \tag{31}
\end{equation*}
$$

where $F(S(t))$ is the egg survival coefficient and $F(B(t))$ is the density-dependent term. The model can be reduced to a piecewise form:

$$
F(S(t))= \begin{cases}D_{1} & \text { if } B(t)<=B_{d}  \tag{32}\\ D_{2} & \text { if } B(t)>B_{d}\end{cases}
$$

where $D_{1}$ and $D_{2}$ are density-dependent coefficients for the survival of white bass eggs and stocked hybrid fingerlings
to age $I, B(t)$ is the total Morone sp. biomass, and $B_{d}$ is a pre-determined carrying capacity.

Density-dependent feedback mechanism in the form of negative exponential function was incoporated into the model. The survival of both species to age $I$ was considered to be under some form of density dependence and was fit with the relationship:

Density dependent coefficient

$$
\begin{equation*}
=\text { ADEN }_{\mathrm{i}} * \exp ^{-(B D E N i * T o t a l ~ b i o m a s s)} \tag{33}
\end{equation*}
$$

where $A D E N_{i}, B D E N_{i}$ are arbitrarily determined constants and total biomass is the combined biomass for both species (Table XIX). The constants $A D E N_{i}$ and $B D E N_{i}$ were determined from iterations in which total biomass values were used along with various combinations of $\operatorname{ADEN}_{i}$ and $\operatorname{BDEN}_{i}$. The final constants were chosen from among the group which yielded density dependent coefficients between 0 and 1 and were similar to those reported for other species (Zagar and Orth 1986; Taylor 1981).

Density-Dependent Effects on Growth

Factors affecting growth of adult fishes include forage availability, overcrowding (both inter- and intraspecific), genetics, and environmental suitability. High density in fish populations can result in reduced fecundity (LeCren 1965), reduced growth (Gulland 1983), and higher mortality of early life history stages through predation, starvation, cannibalism, and competition for space and
other resources.
In my study, the effects of density on the growth of the two species were incoporated into the model via a feedback mechanism that enhanced growth at low densities and retarded it at high densities (Table XIX). The proportion of growth, $\mathrm{PG}_{i j}$, achieved at any given total biomass is given by:

$$
P G_{i j}=\left\{\begin{array}{l}
P G^{*} e^{+}+(b l * \text { total biomass })  \tag{34}\\
(P G=0.9 \text { if total biomass }>700 \mathrm{~kg}) \\
P G^{*} \exp ^{+(b l * t o t a l ~ b i o m a s s)} \\
(P G=1.05 \text { if total biomass }<=700 \mathrm{~kg})
\end{array}\right.
$$

Consequently, the proportion of growth, $\mathrm{PG}_{\mathrm{ij}}$, computed under any given population biomass is applied to the instantaneous rate of growth, $G_{i j}$, and the corrected instantaneous rate of growth is provided thus:

$$
\begin{equation*}
\text { New } G_{i j}=G_{i j} * P G_{i j} \tag{35}
\end{equation*}
$$

The corrected rate was used in all computations in the model. PG is constrained to between 0.90 to 1.05 implying that growth can be retarded by about $10 \%$ or enhanced by $5 \%$ as a result of density-dependent effects.

The effects of high density on growth were examined in the field by comparing the annual variations in body lengths and weights. Annual lengths-at-age data were used to compute proportional stock density (Anderson 1976). Proportional stock density (PSD) is an index developed for assessing the quality of sport fisheries and is expressed as the ratio of number of $f i s h$ in the population that are
at least of stock length to the number that are at least of quality length. Quality and stock lengths employed in the simulation were derived from Gabelhouse (1984). Populations with higher PSDs have greater proportions of bigger fish than populations with lower PSDs.

Model Input and Output

Input into the model consists of: (i) number of age classes in each species, (ii) number of years the simulation is to run, (iii) year to commence simulation, (iv) number of hybrid fry to be stocked annually, (v) agespecific fecundity for white bass, (vi) age-specific instantaneous survival rates for each species, (vii) agespecific frequency of each species, (viii) age-specific mean lengths and mean weights for each species, (ix) agespecific instantaneous total, fishing, and natural mortality rates for each species, and (x) age-specific instantaneous rates of growth for each species.

Model output consists of: (i) Age-specific frequency of each species in the population, (ii) age-specific catch, yield, and biomass of each species during each year of simulation, (iii) age-specific mean lengths and weights for each species during each year of simulation, and (iv) proportional stock density (PSD) (Anderson 1976) for each species during each year of simulation. Input codes for variables and parameters used in the model are listed in Table XX.

## Model Implementation

Factors which influence the usefulness of a model include the ease and extent of the distribution of the language in which the program was written, the codes used and their degree of relatedness to the actual variables, program documentation, and system requirements such as computer time and memory allocation requirements. In order to make this model easily accessible to as many potential users as possible, I (i) used a modular approach by isolating key segments into subroutines, (ii) used arrays, thereby making it easier to vary population parameters from one reservoir to another as well as vary the number of age classes (iii) allow the user to modify the densitydependent effects and stochasticity by modifying the appropriate parameters. This model was written in FORTRAN 77 and was run on an $I B M 3081 K$ mainframe computer system at Oklahoma State University.

## Model Validation

As indicated earlier, one of the benefits derived from population modeling is the synthesis of knowledge and information from different facets of the ecosystem into a workable 'package'. If the assumptions made in initial model development are correct, this package allows for the testing of the interactions between different facets of the
system; the ultimate goal of which is to optimize the allocation of resources and the evaluation of alternative management practices.

Model validation is a two-step process involving first, the testing of the various relationships in the model and secondly, evaluating the joint action of these relationships which define the structure of the model. The object of testing a model is to determine whether the model can adequately account for as much of the behavior of the real system as possible. However, lack of data for all sets of conditions in the system is often the reason for modeling in the first place. Such was the case for the white bass and striped bass $x$ white bass hybrid populations in Lake Carl Blackwell. Consequently, formal and rigorous testing of the results of this model could not be undertaken. The results presented here are the types that can be generally obtained from the model. Comparisons with actual data are made when such comparisons enhance the validity of the model. The model should serve as a prototype for building similar two-species models and testing various multispecies reservoir fishery management hypotheses. A flow chart of the program for this model is shown in Appendix C and the program listing is shown in Appendix $D$.

Results

Population size

Fifty-year simulations, constrained by densitydependent mechanisms and subjected to stochastic variations, produced stable populations of white bass and striped bass $x$ white bass hybrids (Table XXI). However, population size fluctuated widely during the early years and continued to fluctuate thereafter due to the stochastic variation and the compensatory responses to high density (Figure ll). The total annual population size (excluding young-of-the-year) ranged from 2704 to 4386 fish. The inverse relationship between growth rate and population size implies that reductions in population density when they are at high levels brings about much greater growth changes than when population densities are low.

Various scenarios were simulated to determine what annual stocking rate of striped bass $x$ white bass hybrid would be required to maintain an equilibrium with the white bass population. The results of one such simulation (Table XXI; Figure ll) indicates that 10,000 hybrid fingerling stocked annually would maintain the population in equilibrium. This simulation as well as others to be described later, rely on the successful natural reproduction of white bass to sustain that population. In this simulation, white bass were initially more abundant in the population than
the hybrids but by the thirteenth year of the simulation the hybrid population had surpassed that of the white bass. In all simulations, the white bass population was less variable than that of the hybrid even though both populations were subjected to the same degree of stochastic perturbations. Wide fluctuations in the hybrid population is especially apparent during simulation years 23 to 36 (Figure 11).

## Biomass

Biomass can be used to express the quality of a fishery since it is a function of both size and number of fish in the population. A balanced fishery (Swingle 1950) is one with an abundance of intermediate size fishes rather than one with too many small or too few large fishes. A carrying capacity of 700 kg of Morone sp. was imposed on the lake hence total biomass fluctuated about that point (Table XXI; Figure 12). The white bass biomass were higher than that of the hybrid during most of the simulation.

## Fecundity

The density-dependent compensatory mechanism built into the model was such that only a given percentage of the theoretical fecundity was realized each year. Such an approach may be realistic since the expected number of eggs, based on standard age-fecundity or weight-fecundity
relationships is often not achieved when resources such as food and space are limiting. Such requirements are assumed to be limiting under density-dependence thus the resulting effect is decrease in white bass fecundity. Annual fecundity as well as model predicted fertility indicate that fecundity declined during the early years of the simulation but increased thereafter (Table XXII; Figure 13). White bass realized only 35 to $47 \%$ of their theoretical fecundity as a result of the density-dependent constraints imposed by the entire Morone sp. populations biomass.

Percent of White bass in the Population

One of the goals of a multispecies fishery that depends on natural reproduction of one of the species for propagation, is the achievement of acceptable and sustainable yield. The goal of my model, which calls for an annual stocking of hybrid fingerlings, is the maintenance of a viable two-species fishery. The model indicated that white bass and hybrid populations can be maintained within a predictable proportion of the total population. White bass populations constituted 48 to $66 \%$ of the total population (excluding larval fish and young-of-the-year) during any given year (Table XXII; Figure 14). The proportion of either species in the population could be varied by the modeler, through suitable choice of parameters, to suit the needs of the fishery.

## Percent Survival of Stocked

Hybrid Fingerling

The density-dependent constraints imposed on hybrid fingerling survival resulted (Table XXII; Figure 15) in only 2 to $5 \%$ annual survival through the 'critical' period of development. Larval survival is the key to successful development of fish populations and has been shown to account for variability in year-class strengths in some reservoir fish populations. These values of survival estimated by the model seem to be low for Lake Carl Blackwell since Kleinholz (1985) estimated that mortality of hybrid fry stocked in Lake Carl Blackwell ranged from 1 to $5 \%$ and that young-of-the-year mortality ranged from 40 to $50 \%$. Kleinholz (1984) concluded that hybrid fry mortality is highest at stocking and is most common in small reservoirs with inadequate forage. Huner (1985) concluded that survival rates in excess of $50 \%$ for pond-raised hybrid fry were good. Hence higher mortality rates than those estimated by Kleinholz might be expected in resevoirs.

## Harvest

Annual harvest estimates obtained from the simulation ranged from 217 to 397 for the white bass and 244 to 387 for the hybrid (Table XXIII; Figure 16). This level of harvest is small compared to those obtained in other reservoirs the size of Lake Carl Blackwell. However, angling for
white bass and hybrids in Lake Carl Blackwell has traditionally resulted in poorer catches than in surrounding reservoirs. One of the assumptions of the model was that annual age-specific fishing mortality rates (Table XV) were constant. Estimated instantaneous fishing mortality rates were higher for the hybrid (0.09 to 0.77) than for the white bass (0.03 to 0.55). On the average, harvest was greater for the hybrid than for the white bass during most of the simulated years and based on the population parameters employed in the modeling, 5 to $10 \%$ of the white bass and 6 to $14 \%$ of the hybrid (ages I-V) are removed by anglers.

Yield

Yield in fisheries is a reflection of both numbers and weight of fish caught. In commercial fisheries it is also a measure of the energetic contribution of the fish to man's nutritional needs and represents that portion of the biomass that is harvested by man. The yield curves of both species were quite similar to the harvest curves. This similarity is not unexpected since constant annual agespecific fishing mortality rates were employed in the modeling and both species had similar age-specific mean weights. The mean annual yields ranged from 110 to 171 kg for the white bass and from 109 to 186 , $k g$ for the hybrid after the model had stabilized (Table XXII; Figure l7). Thus the combined annual yield for the Morone sp. fishery
ranged from 241 to 333 kg .

## Proportional Stock Density

In addition to expressing harvest in terms of weight and numbers, fishing quality has been variously expressed in terms of population indices such as forage/carnivorous (F/C) ratio or balance (Swingle 1950), young/carnivorous (Y/C) ratio (Swingle 1950), proportional stock density (PSD) (Anderson 1976; 1978), and relative stock density (RSD) (Anderson 1980). Proportional stock density, the proportion of quality length fish that are also at least of stock length, has become popular among biologists as a convenient method of expressing population structure. The simulation indicated that both species had annual PSD's ranging from 41 to 58 for the white bass and from 31 to 61 for the hybrid (Table XXIII; Figure 18). The PSD's of 100 which were obtained during the early part of the simulation are not representative since the populations were not in equilibrium at that time.

## Alternative Hybrid Stocking Rates

One of the most important advantages of models is that they facilitate the quick, inexpensive simulation of management scenarios that are impossible to test or implement in the real system. In keeping with the primary objectives of this study, the effects of various hybrid fingerl-
ing stocking rates were simulated. The stocking rate of $10,000 \mathrm{fish}$ was used as a baseline and the behavior of the system under stocking rates of 15,000 and 5,000 fish were examined for stability, yield and other parameters previously discussed. The rationale for testing different annual stocking rates was to determine which rates would produce persistent and satisfactory yield of both species to the fishery at the least cost.

A 50 -year simulation with an annual stocking rate of 15,000 hybrid fingerlings (all other parameters remaining unchanged) indicated that the hybrid population was consistently higher relative to that of the white bass than at the base stocking rate (Table XXIV; Figure 19). The total Morone sp. population size was slightly higher ( $p<0.05$ ) at this stocking rate (mean annual $=3,600 \mathrm{fish}$ ) than at the baseline stocking rate (mean annual=3,100 fish). However, the $50 \%$ increase in hybrid stocking rate also resulted in a corresponding significant decrease in the white bass population ( $p<0.05$ )

Under the 15,000 fingerling stocking rate, the simulated annual total biomass (Table XXIV; Figure 20) was similar ( $p>0.05$ ) to that at the 10,000 stocking rate but that of white bass significantly lower (p<0.05). White bass fecundity was lower ( $p<0.05$ ) at higher total population density than at lower population density but the proportion of white bass eggs which hatched as a result of density dependence remained essentially unchanged (Table XXV; Figure 21). The proportion of white bass in the population was
lower ( $\mathrm{p}<0.05$ ) at high hybrid stocking density than at the baseline rate (Table $X X V$; Figure 22). At the baseline hybrid stocking rate, white bass constituted 48 to $66 \%$ of the entire population but at the higher rate, they made up only 34 to 52\%. This difference indicates that a $50 \%$ increase in hybrid stocking rate does not translate into a corresponding decrease in the proportion of white bass in the population. The proportion of stocked hybrid fingerling that died annually from density-dependent causes remained essentially unchanged (Table XXV; Figure 23) under both stocking rates. The justification for increasing the hybrid stocking rate is to increase fish harvest and yield. To this end, the increase in hybrid population size resulted in higher ( $p<0.0001$ ) harvests (Table XXVI; Figure 24) and yield ( $\mathrm{p}<0.05$ ) (Table XXVI; Figure 25) resulting in an annual mean hybrid harvest of 366 fish weighing 260 kg . Growth of both species were retarded at high total population densities and were enhanced at lower densities by an amount proportional to the total population biomass. The increased hybrid population size might be expected to result in lower hybrid average size as well as lower PSD. Conversely, white bass might be expected to obtain larger average sizes and have higher PSDs as their population size decreased. However, the density-dependent effects on growth operated on the entire Morone population not each species individually. Consequently, differences in PSD between the two groups were not observed (Table XXVI; Figure 26).

A $50 \%$ decrease in hybrid fingerling stocking rate
resulted in a decrease (p<0.05) in the number of hybrids in the total population (Table XXVII; Figure 27) but the survival rate of stocked hybrid fingerling remained unchanged (Table XXVIII; Figure 28). A persistent and sustainable fishery resulted at the lower stocking rate but the contribution of hybrid to the overall fishery was reduced to about $33 \%$ of that of the white bass. The reduced stocking rate did not result in lower overall Morone biomass but did result in a substantial decrease (p<0.000l) in hybrid biomass (Table XXVII; Figure 29).

White bass fecundity increased with decreased stocking of hybrids. However, the percentage of eggs that hatched remained essentially unchanged (Table XXVIII; Figure 30) because of the compensatory feedback mechanism. Decreased hybrid stocking rates resulted in a $16 \%$ increase in the proportion of white bass in the population (Table XXVIII; Figure 31). The simulated annual harvest (Table XXIX; Figure 32) and yield (Table XXIX; Figure 33) of white bass did not increase in proportion to the $50 \%$ stocking decrease of striped bass $x$ white bass hybrid. Also the PSD increased only slightly at the lower stocking density (Table XXIX; Figure 34). In summary, the lower overall Morone sp. population size brought on by reduced hybrid stocking resulted in an increased $P S D$ for both species and a corresponding increase in the quality of the fishery.

## Slot-Length Limits

Various measures have been adopted to ensure adequate representation of fish of desirable sizes within the population. One of these measures is the slot-length limit whereby only fish within certain size-classes are harvested. The effects of restricting fishing to age classes II through IV were tested for both species and the result indicated that significant differences ( $p<0.05$ ) in annual harvests were observed as a result of the size restriction (Table XXX). However, no differences in yield ( $\mathrm{p}>0.05$ ) occurred (Table $x X X)$. The relatively small contributions made by age-classes $I$ and $V$ to the fishery might explain the lack of significant differences in yield. Despite their relatively large numbers, fishes in age-class I have traditionally not contributed much to sport fisheries because of their small sizes. On the other hand, older and larger fishes are in greater demand but their numbers are usually low; hence both age-classes do not significantly contribute to the yield.

## Growth in Length

There were very few changes in lengths-at-age for the white bass (Figure 35) or the hybrid (Figure 36) in the simulations. Compensatory growth occurred when the biomass dropped below 700 kg and compensation was reflected as a $5 \%$
increase in the instantaneous rate of growth. Conversely, a $10 \%$ reduction in the instantaneous rate of growth occurred when biomass exceeded of 700 . In summary, the density of the Morone sp. fluctuated widely but these fluctuations were not reflected in the growth patterns of either species.

## Discussion

When two species share a common limiting resource, one of the species will be excluded from the resource unless the species can subdivide the use of that resource (Gause 1934; Neyman et al. 1956). This subdivision is achieved when each species excludes their competitor from part of their 'fundamental niches'. The exact requirement for stable coexistence is that each species must inhibit its own growth more than it inhibits that of the other species. This requirement is commonly called the Lotka-Volterra model. One property of the Lotka-Volterra model is that total population size of the combined system is larger than that of each individual group or species. This property served as the basis for my model. I assumed that the total population size attained by the Morone sp. fishery was greater than that of each of the two forms. This assumption dictated that the marginal resources 'empty niches' that are not utilized by either form when they were alone in the system would be used to some extent when both forms were in coexistence.

The major assumptions of this model were that densitydependent effects on recruitment and growth as well as random environmental fluctuations were in effect. However, it is widely accepted that the degree to which recruitment depends on density is one of the more important problems of fishery research. The stock-recruitment process appears to be the main mechanism by which fish populations maintain themselves at around the level they do rather than continuously declining or expanding (Cushing 1977). However, there is also evidence (Parrish 1973; Gulland 1983) that recruitment to most fish stocks bears no obvious relation to the abundance of the parent stock. Rather, it is clear that recruitment in those stocks is governed mostly by environmental factors at some early stage in the life history of the species.

In recognition of the possible manifestations of density-dependence and stochastic environmental fluctuations on fish populations in reservoirs, I made the model flexible in order to accommodate situations other than those in which density-dependence and stochastic perturbations are both operative. Thus the model can be used for populations where either density-dependence or stochastic environmental fluctuations are known or suspected to occur as well as for populations where none of the above mechanisms are operative. In order to nullify the effects of density-dependence, lines 98 to 106 of the program (Appendix D) should be modified such that the value of 1.0 is returned for the coefficients of interest. For example,
division of the fecundity coefficient (EGGDEN) by itself (lines 100-103) nullifies density-dependence on recruitment while division of the random number (RANDEV) by itself (line l79) nullifies random fluctuations in the model.

Although a time interval of one year was assumed in this model, it could be modified to simulate population changes on a monthly, seasonal, or on the basis of any other time interval. With those modifications, the model response will undoubtedly better reflect the environmental and biological events that occur in nature. Environmental changes, such as seasonal floods, and biological events, such as incresed population sizes during the spawning season, could be examined in greater detail. However, such models are rather complex and entail collection of large amounts of data.

Mortality and growth are the two main densitydependent effects on fish populations. Mortality will have the greatest effects early in life, particularly the larval and immediately post-larval stages. Conversely, growth tends to become increasingly important as a fish ages. There are density-dependent growth effects early in life, but it is among adult fish that the influence of population density in growth is most apparent. The simplest form of density regulation for a fish population, density-dependent growth of the adults, occurs because they grow in weight by an order of magnitude subsequent to maturation. The density-dependent mortality of early life stages may operate directly through intraspecific competition for a
limited resource such as food or space or indirectly by lengthening developmental time and thereby increasing vulnerability. White bass in Oklahoma spawn from late March to May and because striped bass $x$ white bass hybrids are also stocked in reservoirs at this time there is potential for competition between white bass larvae and those of striped bass $x$ white bass hybrids.

Changes in growth rates and in densities are customarily explained in terms of changes in food supply (Weatherly and Rogers 1978). Beverton and Holt (1957) explain the relationships thus: "the variation of growth with density in fish populations is perhaps the best established of the density-dependent effects ..."

One of the reasons for hybrid introduction was its fast growth rate and voracious appetite for clupeids (Williams 1972). Several. studies have suggested broader and more aggressive use of these resources by the hybrid than by the white bass (Keith 1986, citing K. E. Erickson, (ODWC)). If these suggestions are true (Maughan et al. 1986; Gleason 1982) hybrids should have a greater impact on the white bass than vice versa. My model is designed to give such a result. Striped bass $x$ white bass hybrids have been considered to be serious competitors to some segments of white bass populations (Kleinholz 1985) and have been suggested to have severe impacts on the community structures of some populations (Keith 1986). However, there are also indications that the detrimental effects of hybrid bass , 久opulations on the endemic sport fishes may have been

Overemphasized (Harper and Namminga 1986).
White bass and striped bass $x$ white bass hybrid populations were modeled using different age-specific survival and mortality constants. Keeping rates and parameters constant during simulations is easy and convenient and allows the modeler to simultaneously test the relationships between various components of the system. However, in the real system, the mortality and survival rates vary annually. The population sizes of the two species predicted by the model, reflect density-dependent effects as well as stochastic processes. White bass had slightly higher survival rates of young-of-the-year than hybrids and higher survival was reflected in the higher numbers of white bass in the baseline simulations. Higher survival rates of hybrids might be explained by food habits. Although white bass and striped bass $x$ white bass hybrid have very similar food requirements (Kleinholz 1985), the hybrid has been shown to be the more voracious feeder (Keith 1986). Optimal foraging theory would suggest that competition between white bass and hybrids might force the white bass to accept a broader forage base than the hybrid would be required to accept. The wider niche breadth of the white bass relative to that of the hybrid would however be reflected in the higher levels of variability observed in the hybrid populations since both populations were subjected to the same levels of stochasticity.

The creation of numerous impoundments in Oklahoma has restricted the habitat suitable for spawning white bass;
they must travel upstream to spawn (Trautman 1981). This requirement for stream spawning sites means white bass recruitment is strongly tied to water level fluctuation. In years of poor white bass recruitment, or in systems where natural spawning is sporadic, stocking of hybrids could be used to buffer the loss of the white bass fishery. Stocking strategies for the hybrid would be different in systems where natural reproduction of white bass were adequate and consistent.

The assumption that the hybrid and white bass interact with one another implies that the population structure of one of the species can be deduced once knowledge of the other is available. However, actual knowledge of the community structure of each of these populations is very valuable since it enables additional definition to be added to the model. Population sizes and the associated age structures of reservoir fish populations were obtained in my study in routine fish sampling. However data to be input into the model can be obtained either empirically or estimated from the literature.

In the model, harvest and yield are dependent upon population size and fishing mortality rates. Hence the trend exibited by these two parameters in the simulations parallels those of the population sizes. Although this modeling exercise was made on the assumption that fishing and natural mortality rates were constant from year to year, these factors are rarely constant. The assumption of constant fishing and natural mortality rates allows the
evaluation of the effects of fishing on the population structure but additional insight could be obtained by measurements of these parameters over shorter time intervals and incoporation of these measurements into the model.

Yield is a function of fishing effort as well as growth. In populations with low growth relative to mortality, sustained yields are lower than in expanding populations with younger age-classes. The production (net and gross) from a fishery is a function of the difference between growth and mortality and is higher when growth rate is high relative to mortality. In the real system, mortality is easier to manipulate than growth, and is generally the factor used to increase fish yield and production. In theory, manipulation of mortality allows us to alter the size and age structure of the populations to suit fishery needs.

The white bass fishery is very popular in Oklahoma and th the introduction of hybrids into Oklahoma reservoirs, the popularities of both species have increased. Gilliland (1981) reported that the enthusiastic support for hybrid fishery in Oklahoma is due to the rapid growth of the fish, ease of capture, and large size relative to the white bass.

The harvest and yield curves derived from my model are comparatively lower than those of most small lakes in oklahoma (Glass 1982; Gilliland 1981; Glass and Maughan 1985). In addition, the fishing mortality rates employed in my model were lower than those encountered in other studies (Glass and Maughan 1985). Lower harvest rates than those
estimated by Glass and Maughan (1985) may however be reasonable, because their data came primarily from harvest in a heated effluent canal. predicted returns from the fishery ranged from $0.08-0.14 \mathrm{~kg} / \mathrm{ha}$ for the hybrids and $0.08-0.15 \mathrm{~kg} / \mathrm{ha}$ for the white bass. Axon and Whitehurst (1985) determined from a national creel survey that mean harvest rates for hybrids in 1978 and 1982 were $0.7 \mathrm{~kg} / \mathrm{ha}$ and $3.0 \mathrm{~kg} / \mathrm{ha}$ for mixed fisheries of hybrids and striped bass. The yields predicted by the model probably reflect the low productivity and angling rates prevalent in the reservoir.

In my simulations, yields of both populations tend to reach equilibrium. Higher fishing or natural mortality on any one species can be observed to result in compensatory growth by both species and a predictable level of output from the fishery.

In the real system, the imposition of slot-length limits on one or both species might be instituted to accomplish changes in both yield and harvest of the target population. For example, slot-length limits might be imposed on populations with low PSDs. Using my model, the effects of the restriction on both the community structures and the fisheries could be evaluated. There are currently no creel limits on white bass and striped bass $x$ white bass hybrids in most Oklahoma reservoirs. On the other hand, there are certain size restrictions on the morone sp. fishery in some Oklahoma reservoirs. Consequently, the inability to distinguish accurately white bass from hybrids
makes enforcement of length limits and (or) creel limits difficult in Oklahoma reservoirs which contain two or more of the Morone sp. Use of the model would allow avoidance of the identification problem by enabling the researcher to test the effects of length limits or other regulations on one or both species.

It is generally accepted that fecundity is independent of adult population size (Lecren 1965) since it is thought that population regulating mechanisms act mainly during the first few weeks after hatching. However, annual variations in fecundity have been related to population density. These conflicting interpretations have led to the concept of fecundity and fertility. In my simulations, the fertility of white bass relative to the fecundity fluctuated from 35 to $48 \%$ in response to the density dependent processes. Reduction in fecundity as a result of forage and resource limitation is not uncommon since McFadden et al. (1965) reported that brown trout from infertile streams had lower growth, and fecundity and wydoski and cooper (1966) reported $16 \%$ difference in brown trout fecundity between oductive and unproductive streams.

Factors which affect fecundity can be densitydependent as well as density-independent. Only the densitydependent effects were considered in this model and these factors include age, size, weight, food supply, and age at first spawning. The tenets on which this model was built include the assumptions that forage was limiting and that growth reduction, resulting from food shortage, are trans-
lated into lower fecundity. The compensatory mortality processes which control white bass fecundity are also assumed to control survival of stocked hybrid fingerling during the first few weeks of life. Major density-dependent sources of mortality to fingerlings include cannibalism, predation, and competition for food and space. The overall effects of these processes on the fingerlings result in annual survival rates of 2 to $5 \%$.

The model did not individually consider the densityindependent factors affecting growth and other physiological processes such as temperature, turbidity, water level fluctuations, and wave action. The effects of these actions on fish recruitment processes can be modeled directly (Orth 1977) or indirectly estimated using stochastic processes (Taylor 1981). The second approach was used in my model. Water level fluctuations in Lake Carl Blackwell play a significant role in determining the level of recruitment for largemouth bass (Zweiacker 1972; Summerfelt and Shirley 1978) and Orth (1977) used water level fluctuations to predict year-class-strengths of largemouth bass in Lake Carl Blackwell. Therefore my modeling approach may not accurately assess the importance of these factors in Lake Carl Blackwell.

Many practical applications of single species models assume that the world is deterministic. One example of this assumption in action is in the concept of maximum sustainable yield (MSY) (May 1978). In the actual fishery, management strategies can, however, be significantly
modified by the presence of environmental randomness and unpredictability. By acknowledging the reality of environmental stochasticity, we are forced to consider the dynamics of populations and to consider trade-offs between limits such as MSY and the fluctuations in the yield. Therefore, there seems to be some need to incoporate environmental stochasicity in the absence of real data into management models in fisheries ecology (May et al. 1979).

The degree of stochasticity utilized in this model is set by the user. This approach makes the model flexible enough for use in reservoir systems with different levels of environmental fluctuation. The model could also be adjusted to produce periodic stochastic variations during simulation. This modification would be useful in terms of modeling the effects of drought or flood on fish productivity in reservoirs.

The model developed in my study emphasizes an alternative approach to the management of multispecies fisheries in reservoirs. The future demand for varied recreational fisheries coupled with the diminishing availability of water resources, especially around major population centers, will necessitate management strategies such as those suggested by this model. Kerby et al. (1983) and Massingill et al. (1983) concluded that in view of the high survival, growth, and production capabilities of the hybrid it could be used for commercial aquaculture and in fee-fishing ponds. Attributes of this model such as the response to high densities, typical of most culture situations, would
make it attractive for use in simulating such situations. The use of hybrids such as the striped bass $x$ white bass, walleye $x$ sauger, and muskellunge $x$ northern pike as well as their parental species might be heightened if management strategies suggested by this model are adopted and developed. The model has demonstrated that stable and sustainable populations of two ecologically similar species can be managed simultaneously both as separate stocks and as a joint fishery. Although the model considered five age classes of each species, it can be altered to include any number of age-classes. This model was developed to serve as a guide for managing multispecicies fisheries consisting of a put-and-take species and a naturally reproducing species. Since data for adequate testing of the model are not available, the model can only serve as a prototype upon which future management-oriented two-species models can be based.

## Model Limitations

Applied models are built with the sole objectives of solving specific problems. In the course of model building, problems do arise which were not considered in the initial analysis. Such problems are often in the form of assumptions about the data or processes being modeled. The limitations of this model include lack of long-term data base of both populations. Lack of long-term data makes estimates of parameters less than accurate and precludes testing of the results. Paucity of data on certain aspects
of the population dynamics of these important sport fishes are also highlighted by the model. The identification of these aspects should stimulate more research in those areas. The model also includes assumptions of constant annual age-specific rates of fecundity, growth, survival, and mortality. Obviously these assumptions are not technically correct. Additional empirical information would increase the reliability of the model. Another limitation of my effort is that $I$ assumed an equal number of age-classes in each stock. To overcome this problem, the model has been constructed so that it can be modified to accommodate a different number of age-classes in each species.

Conclusions

The use of models to predict fish yield and population dynamics is becoming an essential component of fisheries management. Models are needed because biological systems, especially warm water fisheries, are very complex and compartmentalization of information into various interacting units enhances understanding. Models are cost effective and require considerably less time to develop than would be required to conduct research in the real systems. Models usually work better when they are simple and describe noninteracting single species populations but since such populations rarely exist in nature, modeling attempts have been made to describe major interactions in some multispecies systems. The model I have developed is designed to
explain the workings of $a$ warm water fishery consisting of two top-level predators. My goal in developing this model was to help fishery biologists answer questions such as those related to harvest, yield, length limits, overcrowding, and random effects on year-class strengths. Interspecific competition and density-dependent control of animal populations as well as stochastic variations in environmental paramerters were the major assumptions upon which the model was built. Density-dependence appears to be a reasonable assumption in this model since significant diet overlap was reported (Kleinholz 1985) for both species in Lake Carl Blackwell. Fluctuations in environmental parameters, especially water level, has been determined to be an important factor controlling largemouth bass recruitment in Lake Carl Blackwell (Summerfelt and Shirley 1978; Zweiacker 1972; orth 1977). If one accepts these assumptions, the model can be used to simulate relevant decision alternatives confronting fishery managers.

The model is presented as a foundation upon which other models can be built. It demonstrates the effects that recruitment or exploitation on one species-stock can have on the other species-stock when both stocks are under density-dependent constraints. It also illustrates the importance of recruitment to population structure as well as the effects of stochastic variation and density-dependence on recruitment. Multispecies fisheries can be managed as different stocks but when the species are potential competitors, as in this case, sound management decisions must
consider the ramifications of species interactions. This model was not validated due to the lack of data on the species considered but it can be modified and the concepts and procedures adapted for use in other fisheries where adequate data for testing, evaluation, and implementation exist.

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APPENDIXES

APPENDIX A
TABLES

Table I. Morphometric Variables Employed in the Multivariate Discrimination of White bass, Striped bass, and Striped bass $x$ White bass hybrids.

VARIABLE

```
Standard Length
Post-Pectoral Length
Post-Pelvic Length
Pre-Dorsal Length
Caudal Peduncle Length
Pre-Pelvic Length
Head Length
Snout Length
Post-Orbital Head Length
Upper Jaw Length
Lower Jaw Length
Base Length of First Dorsal Fin
Pre-Nasal Length
Second Dorsal Spine Length
Second Anal Spine Length
Pectoral Fins Insertion Distance
Pelvic Fins Insertion Distance
Head Width at Pre-Opercle
Head Depth at Occiput
Head Depth at Pupil
Head Depth at Nostril
Head width at Pupil
Nasal to Opercular Spine Distance
Posterior of second Dorsal Fin
    to Posterior of Caudal Peduncle at Lateral Line
Third anal Spine Length
Base Length of Anal Fin
First Dorsal Fin Height
Orbital Length
Base Length of Pectoral Fin
Base Length of Pelvic Fin
Nasal to First Dorsal Fin Distance
Pelvic Fin to Pectoral. Fin Distance
Pectoral Fin to Anterior of Second Dorsal Fin
Pectoral Fin to First Dorsal Fin Distance
Pelvic Fin to First Dorsal Fin Distance
Anterior of Anal Fin to Anterior of First Dorsal Fin
Anterior of Anal Fin to Posterior of First Dorsal Fin
Posterior of Anal Fin to Anterior of First Dorsal Fin
Posterior of Anal Fin to Posterior of First Dorsal Fin
Anterior of Anal Fin to Posterior of Second Dorsal Fin
Posterior of Anal Fin to Posterior of Second Dorsal Fin
Base Length of Second Dorsal Fin
```

Anal Fin Height Pelvic Fin Height pectoral fin Height
Caudal Peduncle Height
Orbit to Preopercle Distance
Preopercle to Opercle Distance
Eye Diameter
Inter-Nasal Distance
Posterior of Second Dorsal Fin to Pectoral Fin Distance
Body Depth at Third Dorsal Spine

Table II. Meristic Variables Counted on White bass, Striped bass, and striped bass $x$ White bass hybrids.

VARIABLE

```
Number of Scales on Lateral Line
Number of Scales Above Lateral Line
Number of Scales Below Lateral Line
Number of Rays on First Dorsal Fin
Number of Spines on Second Dorsal Fin
Number of Rays on Second Dorsal Fin
Number of Rays on Pelvic Fin
Number of Rays on Pectoral Fin
Number of Rays on Anal Fin
Number of Branchiostegal Rays
Number of Upper Gill Rakers
Number of Lower Gill Rakers
Number of Rays on Caudal Fin
```

Table III. Means and Standard Deviations of Morphometric Variables Used in the Multivariate Discrimination of Striped bass, White bass, and Striped bass x White bass Hybrids.

| VARIABLE | SPECIES |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | White Bass |  | HYBRID |  | STRIPED | BASS |
|  | MEAN | SD | MEAN | SD | MEAN | SD |
| Std Len | 140.12 | 79.64 | 169.17 | 17.95 | 269.19 | 53.42 |
| Post-Pect Len | 97.22 | 58.12 | 118.53 | 13.63 | 185.04 | 37.00 |
| Post-Pelv Len | 83.03 | 49.27 | 103.38 | 11.47 | 165.61 | 35.88 |
| Pre-Dor Len | 59.31 | 34.50 | 68.08 | 7.15 | 109.30 | 20.87 |
| Caud Pedun Len | 27.73 | 14.00 | 31.74 | 2.80 | 54.79 | 10.51 |
| Pre-Pelv Len | 53.81 | 30.29 | 63.60 | 7.97 | 99.21 | 18.75 |
| Head Len | 44.77 | 24.29 | 52.49 | 5.14 | 86.42 | 16.48 |
| Snout Len | 11.28 | 6.91 | 13.95 | 1.87 | 23.61 | 5.08 |
| Post-Orb Head Len | 21.07 | 12.20 | 23.93 | 2.64 | 40.99 | 8.39 |
| Upper Jaw Len | 18.01 | 10.05 | 20.85 | 2.06 | 34.22 | 6.92 |
| Lower Jaw Len | 23.86 | 12.92 | 26.73 | 2.50 | 44.40 | 8.90 |
| ```Base Len ``` | 33.06 | 19.26 | 42.17 | 5.81 | 57.43 | 11.98 |
| Pre-Nasal Len | 11.77 | 6.68 | 14.42 | 1.41 | 23.14 | 4.65 |
| 2nd Dor Spine Len | 8.52 | 3.10 | 10.91 | 3.21 | 17.19 | 3.29 |
| 2nd Anal Spine Len | 14.84 | 7.25 | 14.39 | 2.18 | 15.75 | 2.05 |
| Pect Fin Insert Dist | 17.66 | 11.38 | 21.81 | 3.31 | 33.14 | 6.97 |
| Pelv Fin Insert Dist | 11.05 | 6.78 | 11.76 | 1.63 | 16.89 | 3.46 |
| Head Width at Pre-Opercle Head Depth | 21.82 | 13.35 | 25.04 | 2.81 | 37.07 | 7.74 |
| at Occiput | 35.14 | 21.56 | 39.93 | 4.76 | 57.72 | 11.29 |
| Head Depth at Pupil | 24.38 | 13.58 | 26.28 | 2.55 | 39.43 | 7.65 |
| Head Depth at Nostril | 17.23 | 9.03 | 18.33 | 1.82 | 28.33 | 5.67 |
| Head Width at Pupil | 18.01 | 10.84 | 21.02 | 2.41 | 29.69 | 6.62 |
| Nasal-Operc Spine Dist | 33.29 | 17.78 | 39.07 | 3.87 | 64.57 | 12.34 |
| Post of 2nd Dors Fin - Post Caud Pedun Len at Lat Line | 29.52 | 15. 37 | 34.07 | 3.32 | 56.65 | 11.41 |
| 3rd Anal Spine Len | 16.94 | 8.85 | 16.50 | 2.08 | 23.96 | 3.25 |
| Base Len of Anal Fin | 23.04 | 12.93 | 29.89 | 8.37 | 36.19 | 6.63 |
| lst Dor Fin Height | 24.48 | 14.40 | 30.63 | 4.00 | 42.07 | 7.39 |
| Orbital Len Base Len of pect fin | 12.50 6.31 | 5.73 3.98 | 13.95 7.22 | 1.24 0.85 | 19.49 10.82 | 3.14 2.02 |
| Base Len of Pelv Fin | 7.57 | 5.03 | 8.14 | 1.08 | 9.97 | 2.01 |

(Table III Continued)

| Nasal-lst Dor Fin Dist | 47.69 | 27.98 | 53.51 | 5.37 | 85.85 | 16.62 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Pelv Fin-Pect |  |  |  |  |  |  |
| Fin Dist | 9.28 | 5.99 | 12.62 | 2.06 | 17.35 | 3.49 |
| Pect Fin-Ant of |  |  |  |  |  |  |
| 2nd Dor Fin | 63.70 | 29.61 | 65.60 | 8.64 | 93.40 | 18.30 |
| ct Fin-lst |  |  |  |  |  |  |
| Dors Fin Dist | 32.01 | 20.91 | 37.89 | 4.33 | 50.87 | 9.50 |
| Pelv Fin-lst |  |  |  |  |  |  |
| Dors Fin Dist | 41.93 | 26.01 | 51.72 | 6.45 | 68.42 | 12.68 |
| Ant of Anal Fin-Ant |  |  |  |  |  |  |
| Ant of Anal Fin-Post |  |  |  |  |  |  |
|  |  |  |  |  |  |  |
| Post of Anal Fin-Ant |  |  |  |  |  |  |
| of lst Dor Fin | 70.41 | 41.92 | 87.16 | 10.47 | 126.55 | 25.34 |
| Post of Anal Fin-Post |  |  |  |  |  |  |
| Ant of Anal Fin-Post |  |  |  |  |  |  |
| of 2nd Dor Fin | 37.09 | 22.54 | 44.26 | 6.30 | 55.78 | 10.06 |
| Post of Anal Fin-Post |  |  |  |  |  |  |
| of 2 nd Dors Fin | 23.27 | 14.03 | 28.21 | 3.51 | 36.53 | 6.25 |
| Base Len of |  |  |  |  |  |  |
| 2nd Dors Fin | 24.08 | 13.57 | 31.21 | 3.31 | 47.02 | 11.25 |
| Anal Fin Height | 27.85 | 15.70 | 31.61 | 3.81 | 39.44 | 6.79 |
| Pelvic Fin Height | 29.92 | 17.66 | 33.53 | 3.96 | 48.07 | 8.95 |
| Pectoral Fin Height | 26.05 | 16.32 | 27.66 | 3.00 | 44.84 | 8.88 |
| Caud Peduncle Height | 17.89 | 10.59 | 19.43 | 2.79 | 26.56 | 4.73 |
| Orbit-Preoper Dist | 14.89 | 8.69 | 18.21 | 2.03 | 30.00 | 6.41 |
| Preoper-Operc Dist | 16.26 | 8.46 | 18.65 | 2.22 | 28.11 | 5.05 |
| Eye Diameter | 10.01 | 3.83 | 10.32 | 0.66 | 14.38 | 1.78 |
| Internasal Distance | 7.69 | 4.22 | 9.98 | 0.92 | 14.37 | 3.14 |
| Post of 2nd Dor |  |  |  |  |  |  |
| Fin-Pect Fin Dist | 71.56 | 44.31 | 88.99 | 11.21 | 133.75 | 28.54 |
| Body Depth at 3rd |  |  |  |  |  |  |
| Dor Spine | 42.43 | 27.72 | 52.19 | 7.27 | 69.03 | 13.13 |

Table IV. Significant Morphometric Variables and Linear Discriminant Function Coefficients for Separating White bass, Striped bass, and Striped bass $x$ White bass hybrids.

| VARIABLE | FUNCTION | I | FUNCTION II |
| :---: | :---: | :---: | :---: |
| Caudal Peduncle Length | - 25.23 |  | 17.10 |
| Snout Length | -39.85 |  | 5.30 |
| Base Length of First Dorsal Fin | -9.60 |  | -53.45 |
| Base Length of Pelvic Fin | 25.18 |  | -10.71 |
| Pectoral Fin to Second | 66.31 |  |  |
| Pectoral Fin Height | 66.31 14.21 |  | 43.98 49.81 |
| Internasal Distance | -27.44 |  | -45.96 |
| Constant | -35.93 |  | -47.96 |
| Eigen Value | 29.91 |  | 8.58 |
| Variance | 77.71 |  | 22.29 |
| Canonical Correlation | 0.9837 |  | 0.9564 |

Table V. Canonical Variate Scores from the Discriminant Analysis of Morphometric Characters on White bass, Striped bass, and Striped bass $x$ White bass hybrids.
CAN VAR I CAN VAR II

## WHITE BASS

| 9.47 | 2.19 |
| ---: | ---: |
| 9.62 | 3.02 |
| 9.31 | 1.02 |
| 7.78 | 1.55 |
| 8.43 | 1.96 |
| 10.29 | 1.15 |
| 8.31 |  |

HYBRID

$$
\begin{array}{r}
1.44 \\
0.33 \\
3.69 \\
1.35 \\
2.06 \\
-0.60
\end{array}
$$

$$
-6.31
$$

$$
-5.91
$$

$$
-4.07
$$

$$
-6.59
$$

$$
-7.29
$$

$$
-6.52
$$

STRIPED BASS

| -3.24 | 1.28 |
| :--- | ---: |
| -4.22 | 0.85 |
| -2.53 | 1.54 |
| -3.11 | 0.81 |
| -4.50 | 1.66 |
| -4.86 | 0.68 |
| -3.54 | -2.05 |
| -3.61 | 0.57 |
| -3.15 | 1.77 |
| -3.17 | 1.73 |
| -2.92 | 1.36 |
| -2.64 | -0.72 |
| -4.41 | 2.38 |
| -5.24 | 1.32 |
| -3.46 | 1.08 |
| -5.46 | 2.66 |

(Table V Continued)

```
-3.83
-3.29
-3.07
-4.24
-5.43
    1.00
1.85
    1.18
-0.52
    1.35
```

Table VI. Group Centroids of The Linear Discriminant Functions of Morphometric Variables for White bass, Striped bass, and Striped bass $x$ White bass hybrids.

| SPECIES | FUNCTION I | FUNCTION II |
| :--- | ---: | ---: |
| White bass | 8.96 |  |
| Hybrid | 1.38 | -6.11 |
| Striped bass | -3.81 | 1.04 |

Table VII. Classification Matrix for the Discriminant Function Analysis of Morphometric variables for White bass, Striped bass, and Striped bass $x$ White bass hybrids.

| SPECIES | PERCENT <br> CORRECT |  | NUMBER CLASSIFIED <br> INTO GROUP |  |
| :--- | :---: | :---: | :---: | :---: |
|  | White <br> bass | Hybrid | Striped <br> bass |  |
|  |  |  | 8 | 0 |
| White bass | 100.00 | 0 | 6 | 0 |
| Hybrid | 100.00 | 0 | 0 | 0 |
| Striped bass | 100.00 | 8 | 6 | 21 |
| Total | 100.00 |  |  |  |

Table VIII. Classification Function Coefficients for White bass, Striped bass, and Striped bass $x$ White bass hybrids, and Using log Transformation of the Variables Listed.

CLASSIFICATION FUNCTIONS

| VARIABLE | White bass | Hybrid | Striped bass |  |
| :--- | ---: | ---: | ---: | ---: |
| Caudal Peduncle Length | -300.36035 | -245.57773 | 7.58869 |  |
| Snout Length | -2926.40112 | -2666.80835 | -2422.22412 |  |
| Base Length of First |  |  |  |  |
| Dorsal Fin | -1236.32837 | -737.22559 | -1069.62012 |  |
| Base Length of Pelv Fin | -453.61182 | -558.96631 | -766.17554 |  |
| Pectoral Fin to Second |  |  |  |  |
| $\quad$ Dorsal Fin Distance | 6003.64063 | 5150.37891 | 5120.94141 |  |
| Pectoral Fin Height | 3447.52490 | 2942.44629 | 3224.96753 |  |
| Internasal Distance | -3738.73706 | -3164.17896 | -3350.47729 |  |
|  |  | -3324.34546 | -2654.37329 | -2792.80615 |

Table IX. Classification Matrix for the Discriminant Analysis of Morphometric variables for Independent samples of White bass, Striped bass, and Striped bass $x$ White bass hybrids from some Oklahoma Reservoirs.

| SPECIES | PERCENT <br> CORRECT |  | NUMBER CLASSIFIED <br> INTO GROUP |  |
| :--- | ---: | ---: | ---: | ---: |
|  |  | White <br> bass | Hybrid | Striped <br> bass |
|  |  |  |  |  |
|  |  | 10 | 2 | 0 |
| White bass | 83.33 | 1 | 11 | 0 |
| Hybrid | 91.67 | 0 | 0 | 10 |
| Striped bass | 100.00 | 11 | 13 | 10 |
| Total | 91.67 |  |  |  |

Table X. Classification Scores of Independent Morone sp. samples from selected Reservoirs in Oklahoma. (Initial ID W=white bass, H=hybrid, S=striped bass, and * indicates misclassification).

| Sample <br> Number | $\underset{\text { ID }}{\substack{\text { Initial } \\ \hline}}$ | Classification Scores |  |  | Source |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | White bass | Hybrid | Striped bass |  |
| 6 | H | 2897.21 | 2905.52 | 2882.11 | Canton |
| 12 | H | 2395.11 | 2441.76 | 2429.13 | Canton |
| 30 | H | 2502.24 | 2561.09 | 2526.51 | Optima |
| 42 | H | 2664.74 | 2698.88 | 2672.42 | Ft Supply |
| 48 | W | 2283.02 | 2352.33 | 2302.00 | Canton |
| 49 | W* | 2963.97 | 2965.59 | 2935.24 | Lake Carl Blackwell |
| 50 | W | 2900.60 | 2892.24 | 2852.15 | Lake Carl Blackwell |
| 54 | H | 2387.43 | 2449.13 | 2438.28 | Canton |
| 66 | H* | 3322.33 | 3265.85 | 3236.69 | Lake Carl Blackwell |
| 72 | H | 2293.38 | 2374.22 | 2359.23 | Optima |
| 90 | W | 3318.17 | 3262.00 | 3226.53 | Ft Supply |
| 96 | H* | 3239.21 | 3212.70 | 3196.02 | Great Salt Plains |
| 102 | W | 3192.29 | 3184.87 | 3189.32 | Lake Carl Blackwell |
| 132 | S | 2702.32 | 2738.90 | 2776.71 | Texoma |
| 138 | S | 2615.38 | 2680.74 | 2696.27 | Texoma |
| 150 | S | 2790.24 | 2826.25 | 2862.48 | Texoma |
| 151 | S | 2658.41 | 2716.91 | 2747.06 | Texoma |

Table XI. Principal Components and Sheared Principal Components of Morphometric Variables for White bass, Striped bass, and Striped bass x White bass hybrids.

| PC I PC II | SHEARED PC II |
| :--- | :--- | :--- |

## WHITE BASS

| -9.484 | 0.528744 | -0.37131 |
| ---: | ---: | ---: |
| -9.334 | 0.501355 | -0.38426 |
| -9.012 | 0.489265 | -0.36587 |
| -8.638 | 0.430311 | -0.38899 |
| -8.744 | 0.457625 | -0.37188 |
| -8.696 | 0.446527 | -0.37843 |
| -12.098 | 0.858304 | -0.29119 |
| -12.256 | 0.841580 | -0.32273 |
|  |  |  |
|  |  |  |
|  |  |  |
| -10.822 |  |  |
| -10.284 | 0.655348 | -0.37201 |
| -10.871 | 0.573694 | -0.40228 |
| -10.913 | 0.638500 | -0.39335 |
| -10.881 | 0.628016 | -0.40776 |
| -10.007 | 0.640886 | -0.39200 |
|  | 0.542992 | -0.40656 |

STSRIPED BAS

| -10.837 | 0.371968 | -0.65463 |
| :--- | :--- | :--- |
| -12.034 | 0.346425 | -0.79305 |
| -11.857 | 0.465040 | -0.65864 |
| -11.193 | 0.403952 | -0.65649 |
| -12.151 | 0.465972 | -0.68549 |
| -12.349 | 0.452464 | -0.71751 |
| -12.250 | 0.462904 | -0.69784 |
| -13.054 | 0.436173 | -0.80031 |
| -11.107 | 0.370680 | -0.68141 |
| -11.907 | 0.435280 | -0.69281 |
| -11.080 | 0.373522 | -0.67597 |
| -10.484 | 0.309862 | -0.68284 |
| -12.074 | 0.432741 | -0.71113 |
| -11.276 | 0.408512 | -0.65979 |
| -12.238 | 0.443347 | -0.71617 |
| -12.213 | 0.412074 | -0.74476 |

(Table XI Continued)

| -12.294 | 0.418861 | -0.74572 |
| :--- | :--- | :--- |
| -11.900 | 0.469803 | -0.65998 |
| -11.668 | 0.411261 | -0.69411 |
| -12.207 | 0.432132 | -0.72431 |
| -11.885 | 0.395072 | -0.73064 |

Table XII. Principal Components and Sheared Principal Components of Morphometric Variables for White bass and Striped bass x White bass hybrids.

|  | PC II | SHEARED PC II |
| :---: | :---: | :---: |
|  | WHITE BASS |  |
| -9.47835 | 0.301442 | -0.00662 |
| -9.32603 | 0.268224 | -0.03486 |
| -9.00412 | 0.283347 | -0.00930 |
| -8.62563 | 0.238411 | -0.04190 |
| -8.73361 | 0.283782 | -0.00008 |
| -8.68569 | 0.286446 | 0.00414 |
|  | HYBRID |  |
| -12.11130 | 0.385777 | -0.00786 |
| -12.25600 | 0.393404 | -0.00526 |
| -10.82460 | 0.110387 | -0.24122 |
| -10.27870 | 0.100694 | -0.23317 |
| -10.86960 | 0.151670 | -0.20143 |
| -10.91100 | 0.103046 | -0.25136 |
| -10.88090 | 0.091252 | -0.26216 |
| -9.99984 | 0.141415 | -0.18343 |

Table XIII. Mean Lengths-at-Age, and Estimated Annual Instantaneous Rates of Growth for White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma, during 1985.

|  | White bass |  | Hybrid |  |
| :--- | :---: | :--- | :--- | :--- |
| Age | Mean <br> length | Growth <br> rate | Mean <br> length | Growth <br> rate |
| I | 153 | 0.549 | 188 | 0.503 |
| II | 265 | 0.234 | 311 | 0.208 |
| II | 335 | 0.121 | 383 | 0.106 |
| V | 378 | 0.067 | 426 | 0.057 |
|  | 404 | 0.000 | 451 | 0.000 |

Table XIV. Constants Obtained from Length-Weight Relationships for white bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma, during 1985 (Numbers in parentheses indicate standard errors).

| $N$ | $\log _{10^{a}}$ | $b$ | $R^{2}$ | Species |  |
| :--- | :--- | :--- | :--- | :--- | :--- |
| 83 | -5.03 | $(0.18)$ | 3.05 | $(0.07)$ | 0.9561 |
| 104 | -5.54 | $(0.12)$ | 3.26 | $(0.05)$ | 0.9791 | Hhite bass

Table XV. Age-specific Instantaneous Total, Fishing, and Natural Mortality Rates for White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma during 1985.

| Age | White bass |  |  | Hybrid |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Total | Fishing | Natural | Total | Fishing | Natural |
| I | 0.55 | 0.03 | 0.52 | 0.69 | 0.09 | 0.60 |
| I I | 0.97 | 0.39 | 0.58 | 1.09 | 0.61 | 0.48 |
| III | 0.73 | 0.40 | 0.33 | 1.22 | 0.71 | 0.51 |
| IV | 0.96 | 0.55 | 0.41 | 1.21 | 0.53 | 0.68 |
| V | 1.21 | 0.39 | 0.82 | 1.61 | 0.77 | 0.84 |

Table XVI. Age-specific Instantantaneous Survival Rates for White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma during 1985.

| Age | White bass | Hybrid |
| :--- | :---: | ---: |
| I | 0.53 | 0.43 |
| II | 0.60 | 0.70 |
| II | 0.44 | 0.56 |
| IV | 0.25 | 0.17 |
|  | 0.00 | 0.00 |

Table XVII. Age-specific Fecundity for White bass in the Washita River arm of Lake Texoma (Baglin 1972).

| Age | Fecundity |
| :--- | :---: |
|  |  |
| I | 0 |
| I I | 0 |
| II | 192436 |
| IV | 263677 |
| V | 334918 |

Table XVIII. Monthly Harvests of White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma from June 1984-August 1985.

| Month | 1984 |  |
| :---: | :---: | :---: |
|  | White bass | Hybrid |
| June | 10 | 9 |
| July | 13 | 8 |
| August | 2 | 12 |
| September | 6 | 15 |
| October | 3 | 8 |
| November | 9 | 9 |
| December | 31 | 29 |
|  | 1985 |  |
| February | 18 | 56 |
| March | 11 | 17 |
| April | 13 | 13 |
| May | 4 | 3 |
| June | 1 | 6 |
| July | 34 | 8 |
| August | 2 | 1 |

Table XIX. Constants Used to Simulate the Effects of Densitydependence on Fecundity, Percent Survival of Stocked hybrid Fingerlings, and Proportion of Growth.**

|  | Biomass (kg) |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  |  | $=700$ |  | 00 |
| Density effect | A | B | A | B |
| Fecundity | 0.70 | $-5.0 \times 10^{-7}$ | 0.95 | $-5.0 \times 10^{-7}$ |
| Hybrid survival | 0.04 | $-5.0 \times 10^{-4}$ | 0.08 | $-5.0 \times 10^{-4}$ |
| Prop of Growth | 0.90 | $-5.0 \times 10^{-7}$ | 1.05 | $-5.0 \times 10^{-7}$ |
| ** Models are of the form |  |  |  |  |
| density effect=A*exp (B*total biomass). |  |  |  |  |

Table XX. Input Data Requirements and Codes for Variables and Parameters Used in Modeling Population Dynamics of white bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma.

| Name | Usage Default | Value |
| :---: | :---: | :---: |
| K | Number of age classes in each species (assume equal number) | None |
| NYR | Number of years simulation is to run | None |
| JYEAR | Year simulation is to start | None |
| HSTOCK | Number of hybrid fingerlings stocked annually | None |
| $\mathrm{WSURV}_{\mathrm{i}}, \mathrm{HSURV}_{i}$ | Age-specific survival rates for white bass and hybrid, respectively | None |
| $\mathrm{FERT}_{\mathrm{i}} \mathrm{FN}_{1}$ | Age specific fecundity for white bass | None |
| $\mathrm{WN}_{\mathrm{i}}, \mathrm{HN}_{\mathrm{i}}$ | Number of white bass and hybrid in each age class respectively | None |
| WLEN $_{i},{ }^{\text {HLEN }}$ i | Age-specific mean lengths for white bass and hybrids respectively | None |
| $\mathrm{WG}_{\mathrm{i}}, \mathrm{HG}_{\mathrm{i}}$ | Age-specific instantaneous mean | None |

## (Table XX Continued)

|  | rates of growth for white bass and hybrid respectively |  |
| :---: | :---: | :---: |
| $\mathrm{WWT}_{i},{ }^{\text {HWT }}$ i | Age-specific mean weights for white bass and hybrid respectively | None |
| $\mathrm{WZ}_{\mathrm{i}}, \mathrm{HZ}_{\mathrm{i}}$ | Age-specific instantaneous total mortality rates for white bass and hybrid respectively | None |
| WFMORT $_{i}$, HFMORTi | Age-specific fishing mortality rates for white bass and hybrid respectively | None |
|  | Age-specific natural mortality rates for white bass and hybrid respectively | None |
| WVIVE,HVIVE | Annual survival rate for white bass from egg to age $I$ and survival of hybrid fingerling to age I | None |
| WAWT, HAWT | Constant a in the relationship: | None |
|  | Weight=a*Length ${ }^{\text {b }}$ for white bass and hybrid respectively |  |
| WBWT, HBWT | Constant $b$ in the relationship: | None |
|  | Weight=a*Length ${ }^{\text {b }}$ for white bass and hybrid respectively |  |
| GENDER | Sex ratio for white bass in the population | 0.5 |
| $\mathrm{A}_{1}, \mathrm{~B}_{1}$ | Constants in the relationship: | None |
|  | ```Egg density=a_* exp-(bl*total biomass) for white bass``` |  |
| ${ }^{\text {A }} 11$ | Constant $A_{1 l}$ in the relationship: | None |
|  | $\begin{aligned} & \text { Egg density=all*exp-(bl*total biomass) } \\ & \text { for white bass } \end{aligned}$ |  |
| $\mathrm{A}_{2}, \mathrm{~B}_{2}$ | Constants in the relationship: | None |
|  | ```Survival=a 2* exp-(b2*total biomass) for stocked hybrid fingerling``` |  |
| $\mathrm{A}_{3}$ | Constants in the relationship: | None |
|  | ```Survival=a3* exp-(b2*total biomass) for stocked hybrid fingerling``` |  |
| $\mathrm{A}_{4}$ | ```Constant in the relationship: Prop. of growth=a4*exp-(bl*total biomass) for white bass and hybrid``` | None |
| $A_{5}$ | Constant in the relationship: pfop. of growth=as*exp-(bl*total biomass) for white bass and hybrid | None |
| RANDEV | Normally distributed random variate with mean=l and user supplied std. dev | None |

## (Table XX Continued)

| WTOT, HTOT | a | None |
| :---: | :---: | :---: |
|  | for white bass and hybrid respectively |  |
| TOT | Total population size (both species) | Non |
| WAVLEN $_{\mathrm{i}}, \mathrm{HAVLEN}_{\mathrm{i}}$ | Mean annual lengths-at-age for white bass and hybrid respectively | None |
| $\mathrm{WAVWT}_{i}, \mathrm{HAVWT}_{i}$ | Mean annual weights-at-age for white bass and hybrid respectively | None |
| WBIOMS $_{\mathrm{i}}, \mathrm{HBIOMS}_{\mathrm{i}}$ | Annual age-specific biomass for white bass and hybrid respectively | None |
| WTOTBI, HTOTBI | Total annual biomass for white bass and hybrid respectively | None |
| SUMBIO | Total annual mean biomass (both species) | None |
| WAVBIO $_{i}, \mathrm{HAVBIO}_{i}$ | Mean annual age-specific biomass for white bass and hybrid respectively | Non |
| STKSUV | Density-dependent hybrid fingerling survival rate | None |
| EGGDEN | Density-dependent fecundity coefficient for white bass | None |
| PG | Density-dependent growth coefficient for both species | None |
| $\mathrm{WCATCH}_{\mathrm{i}}, \mathrm{HCATCH}_{\mathrm{i}}$ | Age-specific harvest for white bass and hybrid respectively | None |
| WYIELD $_{i}$, HYIELD ${ }_{i}$ | Age-specific yield for <br> white bass and hybrid respectively | None |
| $W^{\text {WGPROD }}{ }_{i}$, HGPROD $_{i}$ | Age-specific gross production for white bass and hybrid respectively | None |
| $\mathrm{WNEPRO}_{i}$, HNEPRO $_{i}$ | Age-specific net production for white bass and hybrid respectively | Non |
| WSTK, HSTK | Annual number of white bass and hybrid of stock size respectively | None |
| WQUAL, HQUAL | Annual number of white bass and hybrid of quality size respectively | Non |
| WPSD,HPSD | Proportional stock density for white bass and hybrid respectively | None |
| WYLD, HYLD | Total annual yield tor white bass and hybrid respectively | None |
| WHAVST, HHAVST | Total annual harvest for white bass and hybrid respectively | Non |
| YOYCF | Random effects on white bass egg survival and hybrid fingerling survival | None |
| HATCH | 'Fertility' or observed fecundity of white bass due to density-dependence | None |
| SUMEGG | Total theoretical white bass fecundity | None |
| WRATIO | Annual percentage of reproductive potential achieved by white bass | None |
| PERCEN | Annual percentage of adult | None |

## (Table XX Continued)

| HYBFRY | Density-dependent annual hybrid <br> fingerling survival percentage | None |
| :--- | :--- | :--- |
| CLUTCHi | Age-specific total annual | None |
| FAMILY | white bass fecundity |  |
|  | White bass fertility | None |

Table XXI. Simulated Annual Population Size and Biomass (kg) of White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma.

(Table XXI Continued)

| 25 | 3908 | 1984 | 1924 | 816 | 418 | 398 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 26 | 3394 | 1880 | 1514 | 762 | 417 | 345 |
| 27 | 3019 | 1751 | 1268 | 728 | 408 | 320 |
| 28 | 3146 | 1859 | 1287 | 693 | 399 | 294 |
| 29 | 3716 | 1982 | 1733 | 747 | 418 | 329 |
| 30 | 3049 | 1740 | 1309 | 689 | 403 | 286 |
| 31 | 4112 | 2149 | 1963 | 825 | 434 | 391 |
| 32 | 3445 | 1953 | 1491 | 766 | 433 | 333 |
| 33 | 3511 | 2099 | 1412 | 783 | 450 | 333 |
| 34 | 3242 | 1983 | 1259 | 727 | 436 | 291 |
| 35 | 3547 | 2216 | 1331 | 739 | 459 | 280 |
| 36 | 3331 | 2088 | 1243 | 713 | 453 | 259 |
| 37 | 3166 | 1997 | 1169 | 703 | 447 | 256 |
| 38 | 3363 | 2107 | 1256 | 702 | 441 | 261 |
| 39 | 3681 | 2287 | 1394 | 736 | 464 | 272 |
| 40 | 3177 | 1999 | 1179 | 707 | 453 | 255 |
| 41 | 3459 | 2185 | 1274 | 726 | 457 | 270 |
| 42 | 3276 | 2066 | 1210 | 699 | 445 | 254 |
| 43 | 4355 | 2478 | 1877 | 812 | 485 | 328 |
| 44 | 3640 | 2214 | 1426 | 780 | 483 | 297 |
| 45 | 3709 | 2340 | 1369 | 814 | 497 | 317 |
| 46 | 3350 | 2161 | 1189 | 748 | 472 | 276 |
| 47 | 3365 | 2211 | 1154 | 718 | 471 | 247 |
| 48 | 3468 | 2249 | 1219 | 705 | 466 | 239 |
| 49 | 3272 | 2107 | 1164 | 700 | 461 | 2399 |
| 50 | 4386 | 2536 | 1851 | 812 | 491 | 321 |
| 51 | 3838 | 2368 | 1470 | 802 | 503 | 299 |

Table XXII. Simulated Annual White bass Fecundity, Fertility, Percentage of Egg Hatch, Percentage of White bass in the Population, and Percent Survival of Hybrid Fingerlings in Lake Carl Blackwell, Oklahoma.

| YEAR | WHITE BASS |  |  |  | HYBRID |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | FECUNDITY | FERTILITY | \% HATCH | \%WHITE BASS | \%SURV. |
| 00 | 0 | 0 | 0.000 | 100.000 | 4.946 |
| 01 | 305972992 | 107041200 | 34.984 | 55.269 | 1.914 |
| 02 | 251389616 | 87945712 | 34.984 | 66.512 | 1.911 |

(Table XXII Continued)
$\left.\begin{array}{lrrrrr} & & & & \\ 03 & 260379200 & 91092368 & 34.984 & 64.015 & 1.970 \\ 04 & 234123376 & 81908208 & 34.985 & 63.644 & 2.022 \\ 05 & 221948320 & 77653248 & 34.987 & 64.168 & 2.217 \\ 06 & 216414304 & 75718176 & 34.988 & 62.354 & 2.270 \\ 07 & 203563680 & 71222192 & 34.988 & 60.211 & 2.278 \\ 08 & 206355792 & 97986624 & 47.484 & 58.818 & 4.716 \\ 09 & 189887088 & 66433744 & 34.986 & 50.813 & 2.102 \\ 10 & 208297104 & 72877600 & 34.987 & 54.308 & 2.243 \\ 11 & 198275328 & 69371952 & 34.988 & 55.138 & 2.280 \\ 12 & 183943776 & 87345136 & 47.485 & 55.301 & 4.779 \\ 13 & 174237648 & 60960320 & 34.987 & 47.990 & 2.197 \\ 14 & 195556896 & 68420912 & 34.988 & 51.660 & 2.284 \\ 15 & 196643615 & 68800880 & 34.988 & 52.797 & 2.271 \\ 16 & 187536544 & 89049984 & 47.484 & 54.012 & 4.680 \\ 17 & 183313104 & 64136128 & 34.987 & 48.605 & 2.230 \\ 18 & 197931360 & 93985584 & 47.484 & 52.341 & 4.657 \\ 19 & 193355616 & 67646384 & 34.985 & 47.703 & 2.060 \\ 20 & 222113600 & 77710704 & 34.987 & 52.553 & 2.201 \\ 21 & 208362112 & 72899552 & 34.987 & 55.715 & 2.203 \\ 22 & 211478928 & 100418432 & 47.484 & 56.989 & 4.652 \\ 23 & 204221552 & 71450928 & 34.987 & 51.733 & 2.205 \\ 24 & 222462000 & 105633152 & 47.484 & 55.239 & 4.618 \\ 25 & 211718176 & 74071104 & 34.986 & 50.774 & 2.082 \\ 26 & 236009760 & 82571936 & 34.987 & 55.392 & 2.175 \\ 27 & 232753600 & 81434096 & 34.987 & 57.999 & 2.234 \\ 28 & 216854880 & 102970352 & 47.484 & 59.104 & 4.595 \\ 29 & 222888768 & 77981904 & 34.987 & 53.350 & 2.200 \\ 30 & 238428368 & 113214416 & 47.484 & 57.058 & 4.608 \\ 31 & 219416064 & 76763920 & 34.986 & 52.263 & 2.067 \\ 32 & 252922064 & 88488784 & 34.987 & 56.708 & 2.167 \\ 33 & 243687344 & 85257152 & 34.986 & 59.771 & 2.138 \\ 34 & 254165152 & 88925456 & 34.987 & 61.167 & 2.236 \\ 35 & 244440752 & 85522656 & 34.987 & 62.466 & 2.215 \\ 36 & 264646896 & 92593392 & 34.988 & 62.696 & 2.262 \\ 37 & 259008848 & 90621216 & 34.988 & 63.080 & 2.280 \\ 38 & 245845600 & 86015728 & 34.988 & 62.643 & 2.281 \\ 39 & 252994624 & 88515520 & 34.987 & 62.140 & 2.220 \\ 40 & 275077632 & 96243104 & 34.988 & 62.899 & 2.271 \\ 41 . & 252672096 & 88403088 & 34.987 & 63.177 & 2.237 \\ 42 & 261641296 & 124236176 & 47.483 & 63.074 & 4.574 \\ 43 & 253219856 & 88590928 & 34.986 & 56.905 & 2.089 \\ 44 & 294090496 & 102891488 & 34.986 & 60.830 & 2.142 \\ 45 & 279260672 & 97701440 & 34.986 & 63.087 & 2.086 \\ 46 & 283347712 & 99134592 & 344.987 & 64.518 & 2.198 \\ 47 & 266299696 & 93171408 & 344.987 & 65.707 & 2.252 \\ 48 & 266815376 & 93352448 & 34.988 & 64.838 & 2.275 \\ 49 & 273325824 & 129784304 & 47.483 & 64.411 & 4.570 \\ 50 & 260396080 & 91101616 & 34.986 & 57.806 & 2.089 \\ 51 & 299966720 & 104946240 & 34.986 & 61.698 & 2.106 \\ & & & & & \end{array}\right]$

Table XXIII. Simulated Annual Harvest, Yield (kg), and Proportional Stock Density (PSD) of White bass and Striped bass $x$ White Jass hybrids in Lake Carl Blackwell, Oklahoma.

| YEAR | WHITE BASS |  |  | HYBRID |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | HARVEST | YIELD | PSD | HARVEST | YIELD | PSD |
| 00 | 69 | 18 | 100 | 0 | 0 | 0 |
| 01 | 397 | 205 | 100 | 84 | 36 | 100 |
| 02 | 298 | 173 | 100 | 239 | 99 | 100 |
| 03 | 325 | 184 | 100 | 273 | 163 | 100 |
| 04 | 271 | 155 | 100 | 239 | 141 | 100 |
| 05 | 277 | 146 | 100 | 209 | 124 | 100 |
| 06 | 260 | 141 | 100 | 204 | 109 | 100 |
| 07 | 250 | 132 | 100 | 222 | 113 | 100 |
| 08 | 250 | 130 | 100 | 237 | 121 | 51 |
| 09 | 235 | 123 | 100 | 274 | 136 | 100 |
| 10 | 256 | 130 | 55 | 327 | 155 | 59 |
| 11 | 238 | 123 | 54 | 301 | 169 | 57 |
| 12 | 223 | 11.5 | 52 | 253 | 138 | 52 |
| 13 | 217 | 110 | 43 | 277 | 134 | 29 |
| 14 | 243 | 119 | 53 | 348 | 156 | 55 |
| 15 | 239 | 120 | 54 | 335 | 180 | 57 |
| 16 | 229 | 115 | 50 | 286 | 152 | 52 |
| 17 | 227 | 113 | 46 | 293 | 143 | 33 |
| 18 | 244 | 119 | 54 | 339 | 154 | 57 |
| 19 | 243 | 119 | 43 | 365 | 179 | 36 |
| 20 | 273 | 132 | 57 | 387 | 183 | 64 |
| 21 | 254 | 126 | 48 | 332 | 186 | 55 |
| 22 | 260 | 127 | 53 | 279 | 144 | 53 |
| 23 | 253 | 123 | 45 | 289 | 138 | 33 |
| 24 | 274 | 131 | 56 | 335 | 149 | 58 |
| 25 | 263 | 128 | 44 | 351 | 172 | 37 |
| 26 | 293 | 140 | 53 | 370 | 171 | 60 |
| 27 | 280 | 138 | 55 | 322 | 176 | 59 |
| 28 | 267 | 130 | 47 | 269 | 138 | 48 |
| 29 | 277 | 133 | 47 | 289 | 132 | 33 |
| 30 | 290 | 139 | 58 | 333 | 148 | 61 |
| 31 | 275 | 133 | 41 | 346 | 167 | 35 |
| 32 | 313 | 147 | 55 | 370 | 168 | 61 |
| 33 | 299 | 144 | 48 | 331 | 177 | 53 |
| 34 | 313 | 149 | 53 | 286 | 143 | 55 |
| 35 | 302 | 145 | 45 | 266 | 133 | 43 |
| 36 | 327 | 153 | 54 | 271 | 125 | 50 |
| 37 |  | 151 | 53 | 265 | 129 | 52 |
| 38 | 303 316 | 145 | 48 | 256 | 122 | 45 |
| 39 | 316 | 148 | 46 | 271 | 123 | 42 |

(Table XXIII Continued)

| 40 | 335 | 157 | 58 | 282 | 131 | 56 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 41 | 310 | 149 | 47 | 267 | 131 | 47 |
| 42 | 321 | 152 | 53 | 260 | 122 | 50 |
| 43 | 318 | 148 | 42 | 301 | 130 | 31 |
| 44 | 364 | 167 | 57 | 360 | 153 | 61 |
| 45 | 340 | 163 | 49 | 328 | 170 | 55 |
| 46 | 346 | 165 | 54 | 275 | 139 | 56 |
| 47 | 323 | 155 | 49 | 242 | 123 | 47 |
| 48 | 331 | 155 | 49 | 244 | 110 | 44 |
| 49 | 332 | 157 | 54 | 256 | 118 | 50 |
| 50 | 328 | 153 | 42 | 297 | 127 | 31 |
| 51 | 372 | 171 | 54 | 358 | 151 | 58 |

Table XXIV. Simulated Annual Population Size and Biomass (kg) of White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma. (Output with $50 \%$ increase in Hybrid Fingerling Stocking Rate).

| YEAR | POPULATION SIZE |  |  | BIOMASS |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | TOTAL | WHITE BASS | HYBRID | TOTAL | WHITE BASS | HYBRID |
| 00 | 3000 | 3000 | 0 | 601 | 601 | 0 |
| 01 | 3520 | 1590 | 1930 | 1119 | 527 | 592 |
| 02 | 3625 | 2130 | 1495 | 1034 | 640 | 394 |
| 03 | 3493 | 1955 | 1538 | 1029 | 542 | 487 |
| 04 | 3297 | 1850 | 1447 | 976 | 529 | 447 |
| 05 | 3050 | 1737 | 1313 | 838 | 458 | 380 |
| 06 | 3063 | 1676 | 1387 | 804 | 444 | 360 |
| 07 | 3270 | 1710 | 1560 | 808 | 425 | 383 |
| 08 | 3006 | 1528 | 1478 | 762 | 393 | 368 |
| 09 | 3038 | 1496 | 1542 | 752 | 371 | 381 |
| 10 | 3061 | 1455 | 1606 | 730 | 350 | 380 |
| 11 | 2854 | 1322 | 1533 | 689 | 326 | 363 |
| 12 | 3945 | 1497 | 2447 | 846 | 337 | 509 |
| 13 | 3314 | 1370 | 1944 | 772 | 327 | 446 |
| 14 | 3449 | 1459 | 1990 | 818 | 332 | 486 |
| 15 | 3105 | 1345 | 1760 | 747 | 316 | 431 |
| 16 | 3064 | 1354 | 1709 | 709 | 311 | 398 |
| 17 | 2922 | 1266 | 1656 | 661 | 292 | 369 |

(Table XXIV Continued)

| 18 | 4253 | 1507 | 2746 | 832 | 315 | 517 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 19 | 3522 | 1385 | 2137 | 776 | 313 | 463 |
| 20 | 3326 | 1363 | 1963 | 797 | 312 | 485 |
| 21 | 3268 | 1387 | 1880 | 748 | 304 | 444 |
| 22 | 3056 | 1331 | 1725 | 685 | 298 | 386 |
| 23 | 4043 | 1482 | 2561 | 786 | 310 | 476 |
| 24 | 3373 | 1351 | 2022 | 740 | 305 | 434 |
| 25 | 3273 | 1347 | 1925 | 756 | 301 | 456 |
| 26 | 3273 | 1373 | 1900 | 720 | 294 | 426 |
| 27 | 2900 | 1240 | 1659 | 653 | 283 | 370 |
| 28 | 4487 | 1572 | 2915 | 815 | 306 | 509 |
| 29 | 3466 | 1344 | 2122 | 751 | 303 | 448 |
| 30 | 3194 | 1297 | 1898 | 765 | 296 | 469 |
| 31 | 3151 | 1312 | 1839 | 704 | 280 | 424 |
| 32 | 3051 | 1287 | 1764 | 650 | 278 | 372 |
| 33 | 4592 | 1577 | 3014 | 809 | 303 | 506 |
| 34 | 3637 | 1398 | 2239 | 775 | 309 | 466 |
| 35 | 3904 | 1613 | 2291 | 854 | 329 | 525 |
| 36 | 3476 | 1514 | 1962 | 781 | 325 | 456 |
| 38 | 3352 | 1525 | 1828 | 687 | 316 | 371 |
| 39 | 5271 | 1974 | 3297 | 900 | 365 | 535 |
| 40 | 3810 | 1615 | 2195 | 834 | 367 | 467 |
| 41 | 3962 | 1819 | 2143 | 900 | 380 | 520 |
| 42 | 3549 | 1715 | 1835 | 802 | 367 | 435 |
| 43 | 3448 | 1746 | 1702 | 740 | 372 | 368 |
| 44 | 3351 | 1670 | 1682 | 697 | 357 | 340 |
| 45 | 4812 | 2020 | 2792 | 861 | 391 | 470 |
| 46 | 3793 | 1751 | 2042 | 812 | 387 | 425 |
| 47 | 3813 | 1843 | 1969 | 851 | 392 | 459 |
| 48 | 3748 | 1871 | 1878 | 802 | 386 | 416 |
| 49 | 3402 | 1753 | 1648 | 741 | 384 | 357 |
| 50 | 3504 | 1784 | 1720 | 724 | 375 | 348 |
| 51 | 3607 | 1798 | 1809 | 734 | 376 | 358 |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |

Table XXV. Simulated Annual White bass Fecundity, Fertility, Percentage of Egg Hatch, Percentage of White bass in the Population, and Percent Survival of Hybrid Fingerlings in Lake Carl Blackwell, Oklahoma. (Output with $50 \%$ Increase in Hybrid Fingerling Stocking Rate).

(Table XXV Continued)

| 37 | 186982832 | 65420752 | 34.988 | 46.121 | 2.267 |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 38 | 177976640 | 84509856 | 47.484 | 45.482 | 4.618 |
| 39 | 183388592 | 64157120 | 34.984 | 37.450 | 1.947 |
| 40 | 231398800 | 80955776 | 34.985 | 42.391 | 2.052 |
| 41 | 209422096 | 73264736 | 34.984 | 45.918 | 1.947 |
| 42 | 218232288 | 76350656 | 34.986 | 48.314 | 2.106 |
| 43 | 209620944 | 73340176 | 34.987 | 50.636 | 2.213 |
| 44 | 211665664 | 100506112 | 47.483 | 49.821 | 4.579 |
| 45 | 205729536 | 71974304 | 34.985 | 41.976 | 2.008 |
| 46 | 238456640 | 83425936 | 34.986 | 46.163 | 2.090 |
| 47 | 221678512 | 77554432 | 34.985 | 48.347 | 2.025 |
| 48 | 223065264 | 78041520 | 34.986 | 49.903 | 2.106 |
| 49 | 226336816 | 79188544 | 34.987 | 51.544 | 2.212 |
| 50 | 216292896 | 75675104 | 34.987 | 50.910 | 2.242 |
| 51 | 216856384 | 75871856 | 34.987 | 49.844 | 2.224 |

Table XXVI. Simulated Annual Harvest, Yield (kg), and Proportional Stock Density (PSD) of White bass and Striped bass x White bass hybrids in Lake Carl Blackwell, Oklahoma. (Output with $50 \%$ Increase in Hybrid Fingerling Stocking Rate).

| YEAR | WHITE BASS |  |  | HYBRID |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | HARVEST | YIELD | PSD | HARVEST | YIELD | PSD |
| 00 | 69 | 18 | 100 | 0 | 0 | 0 |
| 01 | 397 | 205 | 100 | 125 | 53 | 100 |
| 02 | 298 | 173 | 100 | 352 | 146 | 100 |
| 03 | 325 | 184 | 100 | 388 | 236 | 100 |
| 04 | 271 | 155 | 100 | 330 | 195 | 100 |
| 05 | 277 | 146 | 100 | 283 | 170 | 100 |
| 06 | 260 | 141 | 100 | 278 | 148 | 100 |
| 07 | 250 | 131 | 100 | 305 | 155 | 100 |
| 08 | 250 | 130 | 100 | 328 | 167 | 51 |
| 09 | 229 | 120 | 51 | 327 | 172 | 47 |
| 10 | 222 | 114 | 51 | 330 | 167 | 46 |
| 11 | 213 | 109 | 55 | 335 | 169 | 50 |
| 12 | 202 | 102 | 44 | 390 | 183 | 31 |
| 13 | 219 | 108 | 55 | 473 | 216 | 58 |
| 14 | 209 | 104 | 48 | 452 | 242 | 51 |
| 15 | 216 | 107 | 55 | 403 | 208 | 55 |

(Table XXVI Continued)

| 16 |  |  |  |  |  |  |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 17 | 202 | 101 | 50 | 365 | 192 | 48 |
| 18 | 200 | 99 | 54 | 354 | 172 | 49 |
| 19 | 195 | 95 | 42 | 424 | 188 | 29 |
| 20 | 220 | 105 | 55 | 528 | 230 | 59 |
| 21 | 210 | 103 | 52 | 488 | 260 | 58 |
| 22 | 206 | 101 | 49 | 409 | 212 | 52 |
| 23 | 204 | 99 | 52 | 371 | 189 | 49 |
| 24 | 203 | 97 | 45 | 419 | 185 | 32 |
| 25 | 218 | 104 | 56 | 497 | 219 | 59 |
| 26 | 204 | 100 | 51 | 460 | 240 | 55 |
| 27 | 202 | 98 | 49 | 403 | 202 | 50 |
| 28 | 195 | 97 | 56 | 374 | 187 | 52 |
| 29 | 227 | 92 | 40 | 437 | 187 | 27 |
| 30 | 204 | 106 | 59 | 546 | 231 | 63 |
| 31 | 196 | 95 | 53 | 49 | 395 | 262 |
| 32 | 192 | 92 | 51 | 363 | 180 | 60 |
| 33 | 199 | 92 | 41 | 453 | 184 | 41 |
| 34 | 231 | 107 | 58 | 576 | 242 | 28 |
| 35 | 219 | 104 | 45 | 533 | 277 | 52 |
| 36 | 238 | 112 | 54 | 456 | 228 | 57 |
| 37 | 225 | 108 | 53 | 389 | 204 | 53 |
| 38 | 220 | 105 | 48 | 364 | 172 | 44 |
| 39 | 236 | 108 | 39 | 476 | 189 | 26 |
| 40 | 284 | 130 | 63 | 611 | 254 | 69 |
| 41 | 256 | 123 | 46 | 531 | 287 | 57 |
| 42 | 269 | 127 | 53 | 420 | 216 | 57 |
| 43 | 255 | 121 | 50 | 361 | 187 | 47 |
| 44 | 261 | 122 | 53 | 354 | 162 | 48 |
| 45 | 258 | 121 | 42 | 432 | 180 | 29 |
| 46 | 293 | 135 | 58 | 529 | 223 | 63 |
| 47 | 270 | 129 | 49 | 477 | 249 | 55 |
| 48 | 276 | 130 | 49 | 40 | 51 |  |
| 49 | 274 | 130 | 54 | 367 | 183 | 51 |
| 50 | 265 | 126 | 50 | 356 | 166 | 46 |
| 51 | 267 | 126 | 50 | 369 | 168 | 45 |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |

Table XXVII. Simulated Annual Population Size and Biomass (kg) of White bass and striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma. (Output with $50 \%$ Decrease in Hybrid Fingerling Stocking Rate).

| YEAR | POPULATION SIZE |  |  | BIOMASS |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | TOTAL | WHITE | HYBRID | TOTAL | WHITE | HYBRID |
| 00 | 3000 | 3000 | 0 | 601 | 601 | 0 |
| 01 | 2233 | 1590 | 643 | 724 | 527 | 197 |
| 02 | 2711 | 2130 | 581 | 794 | 640 | 154 |
| 03 | 2550 | 1955 | 595 | 725 | 542 | 183 |
| 04 | 2435 | 1850 | 585 | 707 | 530 | 177 |
| 05 | 2281 | 1737 | 544 | 614 | 458 | 156 |
| 06 | 2834 | 1960 | 873 | 729 | 506 | 223 |
| 07 | 2581 | 1860 | 720 | 670 | 484 | 186 |
| 08 | 2904 | 1986 | 919 | 761 | 512 | 248 |
| 09 | 2524 | 1814 | 710 | 676 | 478 | 198 |
| 10 | 3100 | 2137 | 963 | 779 | 526 | 253 |
| 11 | 2503 | 1812 | 691 | 667 | 480 | 187 |
| 12 | 3048 | 2120 | 927 | 764 | 522 | 242 |
| 13 | 2653 | 1932 | 721 | 680 | 490 | 190 |
| 14 | 3499 | 2443 | 1056 | 830 | 570 | 260 |
| 15 | 2852 | 2100 | 751 | 733 | 537 | 196 |
| 16 | 2871 | 2183 | 688 | 732 | 539 | 192 |
| 17 | 2646 | 2036 | 610 | 658 | 497 | 161 |
| 18 | 3352 | 2427 | 925 | 749 | 547 | 202 |
| 19 | 2957 | 2225 | 731 | 701 | 533 | 169 |
| 20 | 2871 | 2193 | 679 | 704 | 529 | 176 |
| 21 | 2896 | 2234 | 662 | 675 | 512 | 163 |
| 22 | 3428 | 2506 | 922 | 743 | 551 | 192 |
| 23 | 2937 | 2235 | 701 | 695 | 534 | 161 |
| 24 | 3603 | 2638 | 964 | 790 | 577 | 213 |
| 25 | 3107 | 2376 | 731 | 734 | 558 | 176 |
| 26 | 3199 | 2511 | 687 | 738 | 567 | 171 |
| 27 | 2848 | 2255 | 593 | 673 | 528 | 144 |
| 28 | 3839 | 2871 | 967 | 776 | 589 | 187 |
| 29 | 3161 | 2452 | 709 | 726 | 571 | 155 |
| 30 | 3002 | 2366 | 636 | 719 | 558 | 161 |
| 31 | 3022 | 2396 | 626 | 674 | 527 | 148 |
| 32 | 3693 | 2764 | 930 | 744 | 570 | 174 |
| 33 | 3416 | 2654 | 762 | 738 | 580 | 158 |
| 34 | 3306 | 2609 | 697 | 753 | 586 | 167 |
| 35 | 3611 | 2896 | 715 | 763 | 602 | 161 |
| 36 | 3361 | 2736 | 625 | 742 | 603 | 139 |
| 37 | 3186 | 2612 | 574 | 715 | 587 | 128 |
| 38 | 3374 | 2756 | 619 | 712 | 583 | 130 |

(Table XXVII Continued)

| 39 | 3684 | 2996 | 688 | 746 | 611 | 135 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 40 | 3199 | 2616 | 583 | 722 | 596 | 126 |
| 41 | 3491 | 2861 | 629 | 736 | 601 | 134 |
| 42 | 3305 | 2706 | 599 | 712 | 586 | 126 |
| 43 | 3360 | 2753 | 607 | 715 | 589 | 126 |
| 44 | 3231 | 2639 | 592 | 694 | 570 | 123 |
| 45 | 4140 | 3191 | 948 | 787 | 622 | 165 |
| 46 | 3468 | 2765 | 703 | 762 | 615 | 147 |
| 47 | 3590 | 2914 | 676 | 781 | 623 | 158 |
| 48 | 3611 | 2958 | 653 | 759 | 615 | 144 |
| 49 | 3345 | 2774 | 572 | 734 | 610 | 124 |
| 50 | 3407 | 2822 | 585 | 716 | 596 | 120 |
| 51 | 3453 | 2843 | 610 | 719 | 597 | 122 |

Table XXVIII. Simulated Annual White bass Fecundity, Fertility, Percentage of Egg Hatch, Percentage of White bass in the population, and Percent Survival of Hybrid Fingerlings in Lake Carl Blackwell, OKlahoma. (Output with $50 \%$ Decrease in Hybrid Fingerling Stocking Rate).

| YEAR | WHITE BASS |  |  |  | HYBRID |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | FECUNDITY | FERTILITY | \% HATCH | \%WHITE BASS | \% SURV. |
| 00 | 0 | 0 | 0.000 | 100.000 | 4.946 |
| 01 | 305972992 | 107051760 | 34.987 | 71.192 | 2.241 |
| 02 | 251389616 | 87951424 | 34.986 | 78.580 | 2.119 |
| 03 | 260379200 | 91099664 | 34.987 | 76.656 | 2.239 |
| 04 | 234123376 | 81914192 | 34.988 | 75.972 | 2.272 |
| 05 | 221948320 | 105393072 | 47.485 | 76.156 | 4.896 |
| 06 | 216414304 | 75717392 | 34.987 | 69.185 | 2.233 |
| 07 | 232529136 | 110414320 | 47.484 | 72.086 | 4.681 |
| 08 | 230086688 | 80499680 | 34.987 | 68.368 | 2.176 |
| 09 | 240682560 | 114285552 | 47.484 | 71.877 | 4.658 |
| 10 | 225258432 | 78809712 | 34.986 | 68.929 | 2.145 |
| 11 | 252660944 | 119973920 | 47.484 | 72.383 | 4.692 |
| 12 | 229409424 | 80262592 | 34.987 | 69.567 | 2.170 |
| 13 | 251721792 | 119527200 | 47.484 | 72.811 | 4.645 |
| 14 | 240010032 | 83968640 | 34.985 | 69.827 | 2.059 |
| 15 | 287629312 | 100633360 | 34.987 | 73.658 | 2.226 |
| 16 | 267284272 | 93515264 | 34.987 | 76.042 | 2.228 |

(Table XXVIII Continued)

| 17 | 264690192 | 125686464 | 47.484 | 76.928 | 4.726 |
| :--- | ---: | ---: | ---: | ---: | ---: |
| 18 | 249458592 | 87277808 | 34.987 | 72.398 | 2.198 |
| 19 | 286014976 | 100070112 | 34.988 | 75.264 | 2.282 |
| 20 | 279134976 | 97662816 | 34.988 | 76.360 | 2.277 |
| 21 | 269343232 | 127894832 | 47.484 | 77.140 | 4.661 |
| 22 | 269670656 | 94349648 | 34.987 | 73.112 | 2.208 |
| 23 | 299219712 | 142079792 | 47.483 | 76.118 | 4.589 |
| 24 | 281910272 | 98629584 | 34.986 | 73.230 | 2.126 |
| 25 | 313626624 | 109729008 | 34.987 | 76.476 | 2.223 |
| 26 | 296219136 | 103638432 | 34.987 | 78.509 | 2.217 |
| 27 | 303589632 | 144156464 | 47.484 | 79.193 | 4.671 |
| 28 | 280065792 | 97984992 | 34.986 | 74.803 | 2.151 |
| 29 | 335449344 | 117364624 | 34.987 | 77.557 | 2.238 |
| 30 | 312078592 | 109188224 | 34.987 | 78.811 | 2.250 |
| 31 | 291090944 | 138221440 | 47.484 | 79.289 | 4.665 |
| 32 | 287587840 | 100618304 | 34.987 | 74.828 | 2.206 |
| 33 | 327508736 | 114585728 | 34.987 | 77.683 | 2.216 |
| 34 | 331217152 | 115882272 | 34.987 | 78.927 | 2.190 |
| 35 | 321635584 | 112529472 | 34.987 | 80.201 | 2.172 |
| 36 | 345782528 | 120978976 | 34.987 | 81.394 | 2.209 |
| 37 | 338084352 | 118287232 | 34.987 | 81.977 | 2.258 |
| 38 | 322455040 | 112819056 | 34.988 | 81.669 | 2.262 |
| 39 | 331032832 | 115818224 | 34.987 | 81.335 | 2.201 |
| 40 | 360224256 | 126032864 | 34.987 | 81.789 | 2.244 |
| 41 | 330782976 | 115731440 | 34.987 | 81.967 | 2.221 |
| 42 | 342884864 | 119966960 | 34.988 | 81.888 | 2.263 |
| 43 | 331861248 | 116109904 | 34.987 | 81.945 | 2.258 |
| 44 | 334748160 | 158950080 | 47.483 | 81.679 | 4.593 |
| 45 | 324503808 | 113531600 | 34.986 | 77.095 | 2.131 |
| 46 | 376840192 | 131843712 | 34.987 | 79.741 | 2.174 |
| 47 | 350509568 | 122630368 | 34.986 | 81.162 | 2.142 |
| 48 | 352916224 | 123473792 | 34.987 | 81.927 | 2.180 |
| 49 | 357964544 | 125241584 | 34.987 | 82.905 | 2.223 |
| 50 | 342158848 | 119712688 | 34.987 | 82.828 | 2.255 |
| 51 | 342994176 | 120004752 | 34.987 | 82.334 | 2.250 |
|  |  |  |  |  |  |

Table XXIX. Simulated Annual Harvest, Yield (kg) and Proportional Stock Density (PSD) of White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma. (Output with $50 \%$ Decrease in Hybrid Fingerling Stocking Rate).

| YEAR | WHITE BASS |  |  | HYBRID |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | HARVEST | YIELD | PSD | HARVEST | YIELD | PSD |
| 00 | 69 | 18 | 100 | 0 | 0 | 0 |
| 01 | 397 | 205 | 100 | 42 | 18 | 100 |
| 02 | 298 | 173 | 100 | 122 | 51 | 100 |
| 03 | 325 | 184 | 100 | 145 | 85 | 100 |
| 04 | 271 | 155 | 100 | 131 | 77 | 100 |
| 05 | 277 | 146 | 100 | 117 | 69 | 100 |
| 06 | 267 | 146 | 100 | 135 | 69 | 100 |
| 07 | 287 | 151 | 100 | 170 | 83 | 100 |
| 08 | 284 | 152 | 100 | 180 | 100 | 100 |
| 09 | 294 | 156 | 100 | 178 | 95 | 100 |
| 10 | 280 | 149 | 100 | 173 | 98 | 100 |
| 11 | 309 | 161 | 100 | 179 | 91 | 100 |
| 12 | 283 | 150 | 100 | 172 | 98 | 100 |
| 13 | 311 | 161 | 100 | 176 | 89 | 100 |
| 14 | 302 | 157 | 100 | 182 | 99 | 100 |
| 15 | 356 | 180 | 100 | 197 | 97 | 65 |
| 16 | 323 | 169 | 50 | 169 | 100 | 56 |
| 17 | 324 | 166 | 54 | 137 | 75 | 54 |
| 18 | 311 | 158 | 42 | 145 | 72 | 29 |
| 19 | 355 | 175 | 55 | 177 | 80 | 58 |
| 20 | 338 | 172 | 52 | 166 | 91 | 57 |
| 21 | 331 | 167 | 49 | 143 | 76 | 51 |
| 22 | 336 | 167 | 45 | 152 | 73 | 32 |
| 23 | 367 | 179 | 57 | 176 | 81 | 60 |
| 24 | 351 | 175 | 43 | 180 | 92 | 38 |
| 25 | 386 | 189 | 56 | 183 | 88 | 62 |
| 26 | 361 | 180 | 48 | 160 | 88 | 53 |
| 27 | 371 | 182 | 56 | 137 | 70 | 56 |
| 28 | 352 | 172 | 40 | 148 | 69 | 28 |
| 29 | 414 | 197 | 59 | 180 | 78 | 62 |
| 30 | 373 | 187 | 53 | 162 | 88 | 59 |
| 31 | 357 | 177 | 49 | 133 | 69 | 50 |
| 32 | 359 | 173 | 43 | 145 | 66 | 30 |
| 33 | 408 | 191 | 53 | 181 | 77 | 56 |
| 34 | 404 | 196 | 52 | 172 | 89 | 57 |
| 35 | 400 | 193 | 45 | 150 | 76 | 48 |
| 36 | 424 | 202 | 53 | 139 | 69 | 51 |
| 37 | 410 | 198 | 53 | 130 | 64 | 52 |
| 38 | 397 | 192 | 48 | 125 | 60 | 45 |

(Table XXIX Continued)

| 39 | 413 | 194 | 46 | 133 | 60 | 42 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 40 | 439 | 207 | 58 | 139 | 65 | 56 |
| 41 | 406 | 196 | 47 | 132 | 65 | 47 |
| 42 | 421 | 200 | 53 | 129 | 61 | 50 |
| 43 | 405 | 193 | 50 | 128 | 61 | 47 |
| 44 | 411 | 195 | 53 | 128 | 60 | 49 |
| 45 | 408 | 191 | 42 | 150 | 64 | 30 |
| 46 | 463 | 214 | 59 | 180 | 77 | 62 |
| 47 | 426 | 205 | 49 | 163 | 85 | 55 |
| 48 | 437 | 206 | 49 | 139 | 69 | 50 |
| 49 | 434 | 206 | 54 | 128 | 64 | 51 |
| 51 | 422 | 199 | 50 | 125 | 57 | 46 |

Table xxx. Simulated Annual Harvest, Yield (kg) and Proportional Stock Density (PSD) of White bass and Striped bass $x$ White bass hybrids in Lake Carl Blackwell, Oklahoma. (Output With No Fishing Mortality for Ages $I$ and $V$ of Both Species).

| YEAR | WHITE BASS |  |  | HYBRID |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | HARVEST | YIELD | PSD | HARVEST | YIELD | PSD |
| 00 | 0 | 0 | 100 | 0 | 0 | 0 |
| 01 | 397 | 205 | 100 | 0 | 0 | 100 |
| 02 | 271 | 166 | 100 | 204 | 86 | 100 |
| 03 | 304 | 179 | 100 | 239 | 151 | 100 |
| 04 | 227 | 120 | 100 | 206 | 130 | 100 |
| 05 | 258 | 142 | 100 | 161 | 93 | 100 |
| 06 | 232 | 126 | 100 | 161 | 90 | 100 |
| 07 | 222 | 119 | 100 | 173 | 94 | 100 |
| 08 | 227 | 118 | 100 | 196 | 104 | 51 |
| 09 | 206 | 110 | 100 | 191 | 108 | 100 |
| 10 | 234 | 119 | 55 | 283 | 137 | 59 |
| 11 | 216 | 112 | 54 | 257 | 150 | 57 |
| 12 | 202 | 106 | 52 | 209 | 120 | 52 |
| 13 | 187 | 96 | 43 | 180 | 96 | 29 |
| 14 | 220 | 109 | 53 | 297 | 137 | 55 |
| 15 | 218 | 110 | 54 | 289 | 162 | 57 |
| 16 | 206 | 106 | 50 | 237 | 134 | 52 |
| 17 | 199 | 100 | 46 | 201 | 106 | 33 |
| 18 | 221 | 109 | 54 | 290 | 135 | 57 |
| 19 | 213 | 108 | 43 | 271 | 152 | 36 |
| 20 | 251 | 122 | 57 | 343 | 165 | 64 |

(Table $x x x$ Continued)

| 21 | 226 | 114 | 48 | 276 | 158 | 55 |
| :--- | ---: | :--- | :--- | :--- | ---: | ---: |
| 22 | 236 | 117 | 53 | 232 | 126 | 53 |
| 23 | 221 | 109 | 45 | 197 | 102 | 33 |
| 24 | 251 | 122 | 56 | 289 | 132 | 59 |
| 25 | 230 | 115 | 44 | 263 | 146 | 37 |
| 26 | 266 | 129 | 53 | 321 | 153 | 60 |
| 27 | 254 | 125 | 55 | 273 | 150 | 59 |
| 28 | 239 | 120 | 47 | 218 | 121 | 49 |
| 29 | 244 | 117 | 55 | 198 | 98 | 50 |
| 30 | 230 | 113 | 46 | 199 | 102 | 33 |
| 31 | 249 | 120 | 53 | 279 | 127 | 55 |
| 32 | 236 | 116 | 43 | 264 | 141 | 36 |
| 33 | 282 | 134 | 53 | 339 | 158 | 60 |
| 34 | 273 | 133 | 52 | 292 | 159 | 58 |
| 35 | 269 | 132 | 46 | 235 | 128 | 48 |
| 36 | 288 | 136 | 53 | 217 | 105 | 51 |
| 37 | 280 | 136 | 53 | 214 | 109 | 52 |
| 38 | 266 | 131 | 40 | 199 | 103 | 29 |
| 39 | 325 | 150 | 51 | 319 | 136 | 55 |
| 40 | 338 | 162 | 57 | 314 | 166 | 64 |
| 41 | 308 | 153 | 47 | 236 | 133 | 52 |
| 42 | 316 | 149 | 53 | 193 | 95 | 50 |
| 43 | 306 | 146 | 50 | 190 | 94 | 46 |
| 44 | 315 | 151 | 53 | 200 | 99 | 48 |
| 45 | 297 | 143 | 49 | 199 | 99 | 46 |
| 46 | 305 | 145 | 54 | 207 | 101 | 51 |
| 47 | 284 | 137 | 49 | 198 | 100 | 46 |
| 48 | 288 | 138 | 49 | 202 | 99 | 45 |
| 49 | 294 | 139 | 54 | 215 | 104 | 51 |
| 50 | 280 | 135 | 50 | 208 | 106 | 47 |
| 51 | 280 | 134 | 50 | 207 | 102 | 46 |

APPENDIX B
FIGURES



Figure 2. Plot of Principal Component II against Principal Component $I$ for White bass, Striped bass, and Striped bass x White bass hybrids


Figure 3. Plot of Principal Component I against log 10 of Standard Length for White bass Striped bass, and Striped bass $x$ White bass hybrids


Figure 4. Plot of Principal Component II against logio of Standard Length for White bass, Striped bass, and striped bass $x$ White bass hybrids


Figure 5. Plot of Sheared Principal Component II against Principal Component $I$ for White bass, Striped bass, and Striped bass $x$ White bass hybrids


CANONICAL VARIATE I
SPECIES $\stackrel{\rightharpoonup}{\circ} \stackrel{H}{2}$ HYRID ㅁㅁ WHITE

Figure 6. Plot of Sheared Principal Component II against Canonical Variate I for White bass, Striped bass, and Striped bass $x$ White bass hybrids


Figure 7. Plot of Sheared Principal Component II against Canonical Variate II for White bass, Striped bass, and Striped bass $x$ White bass hybrids


Figure 8. Plot of Principal Component II against Principal Component $I$ for White bass and Striped bass $x$ White bass hybrids


Figure 9. Plot of Principal Component II against log of Standard Length for White bass and Striped bass $x$ White bass hybrids


Figure 10. Plot of Principal Component II against Principal Component $I$ for White bass and Striped bass $x$ White bass hybrids


Figure ll. Simulated Total, White bass, and Striped bass $x$ White bass hybrids population sizes (Annual Hybrid Stocking Rate is $\mathbf{1 0 , 0 0 0}$ A=TOTAL, B=WHITE BASS, C=HYBRID)


Figure 12. Simulated Total, White bass, and
Striped bass $x$ White bass hybrids Annual
Biomass (Annual Hybrid Stocking Rate is 10,000 $A=T O T A L, B=W H I T E$ BASS, C=HYBRID)



Figure 14. Simulated Annual Percent of White bass in the Population and Percent Survival of White bass to Age $I$ (Annual Hybrid Stocking Rate is 10,000 A=\% WHITE BASS, $B=\%$ WHITE BASS SURV.)


Figure 15. Simulated Annual Survival Rate of
Stocked Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 10,000)


Figure 16. Simulated Annual Harvest of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 10,000 $A=W H I T E$ BASS, $B=H Y B R I D)$


Figure l7. Simulated Annual Yield of White bass and Striped bass $x$ White bass hybrids
(Annual Hybrid Stocking Rate is 10,000 $A=W H I T E$ BASS, B=HYBRID)


Figure 18. Simulated Annual Proportional Stock Density of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 10,000 A=WHITE BASS, $B=H Y B R I D)$


Figure 19. Simulated Total, White bass, and Striped bass $x$ White bass hybrids population sizes (Annual Hybrid Stocking Rate is 15,000 A=TOTAL, B=WHITE BASS, C=HYBRID )


Figure 20. Simulated Total, White bass, and Striped bass $x$ White bass hybrids Annual Biomass (Annual Hybrid Stocking Rate is 15,000 $A=T O T A L, B=W H I T E$ BASS, $C=H Y B R I D)$


Figure 21. Simulated Fecundity and Fertility
Estimates for White bass (Annual
Hybrid Stocking Rate is 15,000 A=FECUNDITY, B=FERTILITY)



Figure 23. Simulated Annual Survival Rate of Stocked Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 15,000)


Figure 24. Simulated Annual Harvest of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 15,000 A=WHITE BASS, B=HYBRID)


Figure 25. Simulated Annual Yield of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 15,000 A=WHITE BASS, B=HYBRID)


Figure 26. Simulated Annual Proportional Stock Density of White bass and Striped
bass $x$ White bass hybrids (Annual Hybrid
Stocking Rate is 15,000 A=WHITE BASS, B=HYBRID)


Figure 27. Simulated Total, White bass, and Striped bass $x$ White bass hybrids population sizes
(Annual Hybrid Stocking Rate is 5,000 A=TOTAL, B=WHITE BASS, C=HYBRID)


Figure 28. Simulated Annual Survival Rate of Stocked Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 5,000)


Figure 29. Simulated Total, White bass, and Striped bass $x$ White bass hybrids Annual Biomass (Annual Hybrid Stocking Rate is 5,000 A=TOTAL, B=WHITE BASS, C=HYBRID)


Figure 30 . Simulated Fecundity and Fertility
Estimates for White bass (Annual Hybrid
Stocking Rate is 5,000 A=FECUNDITY, B=FERTILITY)


Figure 31. Simulated Annual Percent of White bass in the Population and Percent Survival of White bass to Age $I$ (Annual Hybrid Stocking Rate is 5,000 A=\% WHITE BASS, B=\% WHITE BASS SURV.)


Figure 32. Simulated Annual Harvest of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 5,000 A=WHITE BASS, B=HYBRID)


Figure 33. Simulated Annual Yield of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 5,000 A=WHITE BASS, B=HYBRID)


Figure 34. Simulated Annual Proportional Stock Density of White bass and Striped bass $x$ White bass hybrids (Annual Hybrid Stocking
Rate is 5,000 A=WHITE BASS, B=HYBRID)


Figure 35. Simulated Annual Lengths-at-Age for White bass (Annual Hybrid Stocking Rate is 10,000)


Figure 36. Simulated Annual Lengths-at-Age for Striped bass $x$ White bass hybrids (Annual Hybrid Stocking Rate is 10,000)

## APPENDIX C <br> LOGIC ELOW CHART OF PROGRAM FOR SIMULATING THE POPULATION DYNAMICS OF WHITE BASS AND STRIPED BASS X WHITE BASS HYBRIDS IN LAKE CARL BLACKWELL, OKLAHOMA

SUBROUTINE HYBRID(HVIVE)


SUBROUTINE NORMAL(ISEED, AVERAG, STDDEV, RANDEV)


SUBROUTINE RANDOM(ISEED, U)







## APPENDIX D LISTING OF COMPUTER PROGRAM FOR SIMULATING THE POPULATION DYNAMICS OF WHITE BASS AND STRIPED BASS $X$ WHITE BASS HYBRIDS IN LAKE CARL BLACKWELL, OKLAHOMA








```
            7 I-1,K
            HREXP(I)=EXP(HHF(I))
            HEEXP(I)=EXP(HBG(I))
            HEXPG(I)=EXP(HG(I)*PG)
            HAVLEN(I)=HLEN(I)*HEXPG(I
            HAVWT(I) =HWT(I)* HEEXP(I)
            HBIOMS(I)=HN(I)*HWT(I)*O.001
l
    continue
COMPUTE mEAN bIOmASS at age for white bass
            нтотвI=0.0
            WTOTBI=0.0
            DO B I= 1.K
            IF (WBG(I) .LT.WZ(I)) THEN
                WZEXP(I)=-(WZ(I)-WBG(I)
                WZZUPER(I)=1-WZZEXP(I)
                WZUPER(I)=1-WZZEXP(I)
            ELSEIF (WBG(I).GT.WZ(I)) THEN
            WZEXP(I)=(WEG(I)-WZ(I))
            WZZEXP(I)=EXP(WZEXP(I))
            WZUPER(I)=(WZZEXP(I)-1)
                WZDENM(I)=WEG(I)-WZ(I)
            ENDIF
                WAVBIO(I)=WBIOMS(I)*WZUPER(I)/WZDENM(I)
C
    COMPUTE MEAN BIOMASS AT AGE AND TOTAL BIOMASS FOR HYBRIO
            IF (HBG(I).LT. HZ(I)) TMEN
                HZEXP(I)=-(HZ(T)-HBG(I))
                HZZEXP(I)=EXP(HZEXP(I))
            HZUPER(I)=1-HZZEXP(I)
            HZDENM(I)=HZ(I)-MBGII)
            ELSEIF (HBG(I).GT. HZ(I)) THEN
                HZEXP(I)=(HBGGII)-HZ(I))
            HZZEXP(I)=EXP(HZEXP(I))
            M}\begin{array}{l}{\mathrm{ HZUPER(I)=(HZZEXP(I)-1)}}\\{\mathrm{ HZDENM(I)=HBG(I)-HZ(I)}}
            ENDIF
                HAVBIO(I)=1BEIOMS(I)PHZUPER(I)/HZOENM(I)
            MTOTBI=HTOTEI +HAVBIO(I)
            WTOTBI WTGIEI+WAVBIO(I)
            MPUTE TOTAL ANNUAL MEAN BIOMASS
                BIOMAS(I)=HAVBIO(I)+WAVB
    8. SUMBIO=SU
l CONTINUE
```













Maurice Ibechema Muoneke
Candidate for Degree of
Doctor of Philosophy

Thesis: POPULATION DYNAMICS OF WHITE BASS AND STRIPED BASS X WHITE BASS HYBRIDS IN LAKE CARL BLACKWELL, OKLAHOMA

Major Field: Zoology
Biographical:
Personal Data: Born in Orlu, Imo State, Nigeria, January 31, 1954, the son of Luke N. and Grace N. Muoneke. Married to Ada F. Ezie on November 18, 1983.

Education: Graduated from Comprehensive High School, Orlu, Imo State, Nigeria, June 1973; graduated from Federal Government College, Kaduna, Kaduna State, Nigeria, June 1975; received Bachelor of Science degree in Marine Biology from Nicholls State University, Thibodaux, Louisiana, May 1979; received Master of Science degree in Fisheries from Louisiana State University, Baton Rouge, December 1982; completed requirements for the Doctor of Philosophy degree at Oklahoma State University, May 1987.

Professional Experience: Graduate research assistant, Louisiana Cooperative Fish and Wildife Research Unit, Louisiana State University, Baton Rouge, Louisiana, August 1979 to December 1982; Graduate research Assistant, Oklahoma Cooperative Fish and Wildlife research Unit, Oklahoma State University, Stillwater, Oklahoma, January 1983 to June 1986; Graduate Teaching Assistant, Department of Zoology, Oklahoma State University, Stillwater, Oklahoma, August 1986 to present. Research biologist, Texas Parks and Wildife Department, January 1987 to present.

Professional Organizations: American Fisheries SocietyEarly life History Section. Gamma Sigma Delta (honor Society for Agriculture).

