Neal J. Shunk for the degree of Master of Science in Forest Resources presented on October 5, 2000. Title: Choosing Efficient Land Allocations and Forest Management Regimes for Biodiversity.


Increasing rates of species imperilment and the loss of biological diversity in naturally functioning ecosystems can be directly linked to accelerated urban development and the conversion of natural habitats to satisfy the needs of man. In combating this loss of biodiversity, scientists and policy makers alike recognize the relevance of habitat conservation. This research, funded by a cooperative grant with USDA Forest Service, relies on a framework for modeling wildlife diversity presented by Montgomery et al (1999), to reveal cost effective habitat conservation strategies. Building on this earlier model, alternative forest management strategies are introduced: information that is vital to timber-based economies.

196 mammalian, reptilian, amphibian, and avian species were used to construct a biodiversity index relevant to the Muddy Creek Watershed, Benton County, Oregon. This index, comprised of a taxonomic diversity measure (May 1990) and a classic logistic viability function, measured gains in biodiversity scaled against the opportunity costs of reallocating lands to meet conservation goals. These
index values and associated opportunity costs were calculated and reserved across the full range of land allocation possibilities for the watershed, and formed a marginal cost curve for biodiversity.

The wildlife diversity index ranges in value from 296.19, corresponding to a high development market value maximizing solution, to 310.18 at a cost of nearly 460 million dollars, the highest attainable biodiversity index for this watershed. Forest management played an integral role in the conservation of biodiversity, whereby biodiversity maximizing solutions allocated an overwhelming percentage of forested lands to non-harvested forested reserves.

Two supplementary analyses were undertaken. The first tracked changes in the biodiversity index when management strategies targeting imperiled species were specifically optimized. Land allocations favoring these species had drastic implications on the predicted populations of the remaining non-imperiled species, indicative of the need to consider a broader set of species and their related needs in future land management planning efforts. The second, examined the efficiency of the Institute for a Sustainable Environment's (ISE) high conservation land allocation projections for the year 2025, and verified the necessity of biological indicators in land planning.

# Choosing Efficient Land Allocations and Forest Management Regimes for Biodiversity 

## by

Neal J. Shunk

## A THESIS

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## APPROVED:

n
Signature redacted for privacy.
Major Professor, Repredenting Forest Resources

Signature redacted for privacy.
Chafir of Department of Forest Resources

Signature redacted for privacy.
Dean of Graduaté School

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In avoiding an Emmy-like acceptance, I will attempt to be brief.

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## CHOOSING EFFICIENT LAND ALLOCATIONS AND FOREST MANAGEMENT REGIMES FOR BIODIVERSITY

### 1.0 INTRODUCTION

Declining levels of biodiversity is of growing concern. With urbanization rates steadily climbing, the need for understanding how current land allocations and land use decisions impact our ecological diversity is apparent. In October of 1997 more than six hundred world-renowned scientists met in Washington, D.C. to review the current state of biodiversity worldwide. This group estimated that "species are disappearing at more than 1,000 times the normal rate," and that "seventy five percent of the world's species will be extinct by the end of the next century" (Jewett 1997).

These alarming extinction rates have been the focus of debate for decades, and have ultimately led to a stronger emphasis on wildlife conservation in land use and forest planning. Consider for instance the USDA Forest Service policy initiatives spawned from declining populations of the Northern Spotted Owl, and more recent concerns and activities relating to the preservation of wild salmonid species throughout the Pacific Northwest (Independent Multidisciplinary Science Team 1999).

These programs, aimed at restoring viable breeding populations to endangered species, have proven costly over recent decades. Scientists and policy makers alike now recognize the importance of identifying critical habitats for all species before current populations decline to levels associated with endangerment. This broader concern for ecological diversity has both heightened awareness of the surrounding
issues, and generated millions of research dollars applicable to biodiversity studies. The Oregon Biodiversity Project for example, begun in 1994, has spent four years and over a million dollars mapping Oregon's wildlife habitats (Denson 1999).

Projects like these play an integral role in defining future alternatives for the management of biodiversity. Describing current conditions of ecosystems, and biological relationships that exist between species and their habitats is a necessary first step towards defining an efficient framework for managing for biodiversity. Economists have used such information to examine a variety of related questions regarding future land management and associated gains in wildlife diversity, termed recently as 'the calculus of biodiversity' (May 1990).

These economic studies have added to a growing base of literature addressing the tradeoffs associated with societal demands for natural resources and wildlife biodiversity. Authors have examined a range of associated topics from a cost effective reserve site selection that is both spatial (Csuti et al. 1997a) and non-spatial (Faith et al. 1996), to research aimed at identifying the marginal cost of single species viability (Montgomery et al. 1994, Hyde 1989, Haight et al. 2000). "Pricing Biodiversity" (Montgomery et al.1999) addressed efficient land allocation decisions for varying target levels of biodiversity That study laid the foundation upon which I have constructed my own assessment of biodiversity in the Muddy Creek Watershed, Benton County, Oregon. The Muddy Creek Project utilizes Montgomery's framework for modeling the expected diversity of a given area as a function of species viability and uniqueness. Revealing cost effective tradeoffs between market valued uses and the conservation of biodiversity, the study introduces methodologies
that capture the idea of managing forests for both timber production and wildlife habitat.

The project is aimed at defining efficient land use allocations when biodiversity is of concern. As such, the research finds optimal arrangements of favored habitat at the minimal cost. One hundred ninety six wildlife species are evaluated for their contribution to expected biodiversity, and marginal cost curves are presented for a range of associated habitat allocations. In addition, the study contrasts a diversity index dependent on the wildlife diversity depicted by the full range of species examined, and a second representative of management strategies targeting imperiled and critically imperiled species alone. Current forest management regimes, and the impact that changes in these regimes will bring to the expected biodiversity measure, are examined for both indices.

### 1.1 Objectives

The overall objective of this research is the continued development and analysis of tradeoffs associated with land allocation and the conservation of wildlife biodiversity. The study will focus upon the application and further development of earlier models to evaluate the current state and possible futures of biodiversity in the Muddy Creek Watershed.

In defining efficient land allocations and the tradeoffs between market valued and conservation favoring land uses, a marginal cost curve for biodiversity represents the optimal arrangement of habitats for a range of relative values for wildlife diversity. Points along this marginal cost curve represent optimal choices for local
land managers, and represent the full range of efficient choices for the protection of biodiversity in the watershed. Such information is the fundamental objective of this research and corresponds to the following project objectives:

1. Construct a marginal cost curve for biodiversity relative to the Muddy Creek Watershed.
2. Examine shifts in current forest management regimes resulting from an increased willingness to pay for biodiversity.
3. Develop and contrast a diversity index representative of management strategies targeting imperiled species.
4. Examine the "Possible Futures for the Muddy Creek Watershed, Benton County, Oregon" (Hulse et al. 1997) high conservation landscape's contribution to species population indices, and the resulting efficiency of such solutions.

### 1.2 Literature Review

Human encroachment into previously undisturbed or natural areas has had a direct influence on declining levels of biodiversity. Examining the degree man has influenced shrinking native wildlife populations, as well as defining land management alternatives that most efficiently slow these declines, has led to the amalgamation of biological and economic principles.

In analyzing the tradeoffs that exist between the conservation of wildlife diversity and market valued uses, authors have drawn on a variety of disciplines and examined a range of related topics. These studies have yielded information relating
to population viability, diversity measures, efficient multiple use forest management, reserve site selection, and multiple species biodiversity models. Each will be reviewed in turn.

### 1.2.1 Viability Functions

Economic principles applied to the biodiversity dilemma have been fruitful. Stated simply, "Economics matters because human behavior generally, and economic parameters in particular, help to determine the degree of risk to a species" (Shogren et al. 1999). Defining this degree of risk to a species is often viewed conversely as the probability of species survival, with survival rates typically demonstrated as viability functions.

Viability functions approximate the relationship that exists between population size, given as a function of habitat availability and/or species-specific traits, and the probability of species survival. These curves are often specified as logarithmic, capturing the population dynamics and critical densities associated with a species. This represents larger, marginal viability gains for populations of modest but not yet critical size, whereas lesser marginal gains are associated with the contribution of an additional member to an already thriving population. The Allee effect, depicted by the left tail of the viability function, reflects the imperilment of endangered species. Thus, the probability that an offspring successfully reproduces approaches zero as population size diminishes past a critical threshold due to both the lack of available breeding partners and the distance that species may have to travel to find such a mate.

Early viability modeling efforts were typically focused on depicting the ecological and biological characteristics inherent to the survival of a particular species. Armbruster and Lande (1993), for example, examined the population dynamics of African Elephants in the Tsavo National Park, Kenya. Based upon species fecundity, survivorship, and landscape dynamics, elephant viability was forecasted based upon reserve size. Ruggiero et al.(1994), suggested that viability may best be described by evaluating the following six indicators: disjoint or connection of habitat, habitat separation, age class of forested habitats, habitat size, reproduction rates, and environmental conditions affecting carrying capacities.

In evaluating viabilities for multi-species modeling, data relating to habitat structure and individual specific indicators are difficult to obtain. The International Union for Conservation of Nature and Natural Resources (IUCN) is one of many conservation groups globally that assess species imperilment, where species are assigned an imperilment ranking based upon extent of occurrences, area of occupancy, area and quality of habitat, frequency and locations of sub-populations, and number of mature individuals (IUCN 1994). Mace and Lande (1991), and Mace and Stuart (1994), in association with the IUCN, defined the probability of survival over a specified time horizon given a species' IUCN imperilment ranking. Utilizing these probabilities of survival and the estimated population size intrinsic to the ranking system, Montgomery et al. (1999) constructed a viability curve for use in depicting population dynamics for all species that have been classified according to IUCN ranking standards.

### 1.2.2 Diversity Measures

Diversity measures aid in distinguishing priorities for conservation, a goal inherent to modeling biological diversity. Priorities are often influenced by a species’ relative uniqueness, but preference could also be influenced by aesthetic uniqueness, contribution to ecosystem function, species interdependence, or humanistic features such as future medicinal importance, intrinsic value as watchable wildlife, or potential sociological symbolism.

Taxonomic distinctiveness, sometimes used to summarize the contribution of species uniqueness to a biodiversity index, is typically represented by cladistic-based measurements. Although favorable in some circumstances genetic-based distance measures (Krajewski 1994, Faith 1994, Farris 1979) are rare for most species, and thus their practical application in most studies is not feasible. One such example, Solow et al. (1993) utilized a set of pair-wise genetic distances to differentiate diversity in fourteen crane species.

Cladistic-based diversity measures, attributable to May (1990) and VaneWright et al. (1991), define uniqueness based upon weighted (Cousins 1991) or unweighted taxonomic trees. This approach requires less data, a location specific cladogram, and is a system by which species are ranked relative to one another. The methodology develops diversity rankings based on the number of taxonomically related species sharing the same family, order, or genus, and is a common approach to account for species uniqueness in current biodiversity studies.

### 1.2.3 Efficient Multiple Use Forest Management

Research aimed at identifying efficient forest management regimes, and the tradeoffs that exist between timber production and habitat protection, has been ongoing for several years. Such information is crucial to forest planners, who often work under state-level and congressional directives aimed at reducing biological risk stemming from management.

Initially, such research was aimed at identifying cost effective alternative management actions and the implications these actions had on single species viability. Biologists would often act as the catalyst for management, indicating appropriate keystone or indicator species, whereas threatened or endangered species were the general focus of most early modeling efforts. Biodiversity in these frameworks was negatively impacted by declines in an already endangered species' population.

These single species models ordinarily defined efficient landscape allocations in the face of species' endangerment or extinction. Hyde (1989), established one of the first of these single species models, associating the marginal cost of red-cockaded woodpecker habitat preservation with increased viability for the species. In a similar study, Montgomery et al. (1994) defined the marginal cost of northern spotted owl critical habitat relying on population dynamic models (Lamberson et al. 1992). This model linked the probability of owl survival to reductions in annual timber supply from federal forests in the Pacific northwest.

Haight (1995), presented a generalized model for extinction risk and economic costs in forest planning. This stochastic simulation model suggested methodologies relying on common economic maximization techniques and
constrained by minimum allowable population risk, to determine cost effective conservation plans. The model, repeatedly solved for varying levels of risk, is notable in that it attempts to provide a framework by which USDA Forest Service regulations for sensitive species can actually be accounted for in long-term management planning.

As an application of this framework, Haight, (1999) suggested efficient land management solutions for habitat availability for the San Joaquin kit fox. Re-solving this model with incrementally higher upper bounds on funding, the authors suggested optimal habitat protection areas based on the probability of population extinction, size of protection area, and total amount of funding, subject to a budget constraint.

### 1.2.4 Reserve Site Selection

A broader class of algorithms, reserve site selection optimizers, define efficient biological reserve areas given current habitat configurations, costs for acquisition, and species habitat requirements. One such example (Ando et al. 1998), utilized integer-programming techniques to select reserve areas based upon county level data on land prices and the distribution of endangered species within the United States. A similar approach, Csuti et al. (1997a), compared the efficiency of four reserve site selection algorithms for an Oregon specific terrestrial vertebrate data set. The study contrasted the effects of selecting reserves based on species rarity or endangerment, maximizing species uniqueness indicators. This study did not, however, incorporate land value constraints in the optimization process.

An alternative approach (Haight et al. 2000), introduced reserve selection methodologies that maximized the occurrence of vegetative communities subject to a budget constraint. Based upon the uncertainty of species presence or absence data the authors relied on probabilistic integer optimization techniques to forecast efficient site reserves. In this framework, a vegetative community contributed to the reserve site's production potential when its probability of occurrence exceeded minimum reliability thresholds.

### 1.2.5 Multiple Species Biodiversity Models

Multi-species land allocation or management problems, incorporate many of the ideas already discussed relating to population viability analysis, species uniqueness indicators, single-species biodiversity models, and reserve site selection algorithms. They are unique, however, in that they typically indicate the expected gains in multi-species biodiversity based on future land management policy scenarios. This information is key to the ongoing debate defining what biodiversity is, and how our land management decisions today can be reflected in future healthy populations of wildlife species.

White et al. (1997), for example, assessed multiple species risk by devising species abundance measures predicted by species area requirements, frequency of habitat, and patch size. Implications of future biodiversity changes were tracked across predicted future landscapes, defining a ratio of present to future species abundance.

Rather than defining the change in biodiversity for predicted future landscapes, other authors have chosen to take an economic approach to modeling species-habitat relationships, where efficient or optimal landscape patterns that either meet biodiversity objectives at the minimal cost, or track changes in biodiversity for an increased willingness to pay for the commodity. Hof and Raphael (1992) and Bevers et al. (1995) for instance, defined optimal timber age-class distributions and forest cover types respectively, when constrained by population viability requirements for biodiversity.

Montgomery et al. (1999), developed a marginal cost curve for 147 native bird species in the Poconos region of Monroe County, Pennsylvania. This study traced the supply of expected diversity for incremental changes in the willingness to pay for biodiversity, expressed as foregone land value as land uses changed to accommodate biodiversity. The study incorporated measures of species viability and uniqueness (cladistic-based) that uniquely distinguished it from earlier models relying on population maximizing algorithms.

### 1.2.6 Literature Review Conclusion

Currently, modeling efforts are constrained by the type and validity of biological data available. With advancements in technology and computing capabilities, spatial heuristic solutions to landscape problems could present new opportunities to further our understanding of biodiversity management. Current efforts typically optimize spatial habitat configurations, edge, and/or fragmentation, like Shannon's index (Holland 1994).

Scientific data relating species preference and interaction with specific habitat characteristics, however, is lacking for most species. With the further development of such measures for all species, future land allocation solutions for biodiversity, and the information contained therein, will surely add to our further understanding of the preservation of wild populations.

Given the information currently available, however, this project revealed efficient land allocation alternatives, relying on a biodiversity index similar to Montgomery et al. (1999). Based on the watershed's timber reliant economy a number of forest management alternatives were presented, and gains in biodiversity brought about by changes in current management regimes were forecasted. In addition, the implications of land management strategies targeting imperiled species, as well as community group landscape planning efforts were analyzed. The project united earlier biological diversity models and defined efficient habitat protection strategies for multi-species biodiversity.

### 2.0 METHODS

### 2.1 Study Site Description

The Muddy Creek Watershed, Benton County, Oregon, encompasses some 125
square miles ( 32,000 hectares) in the Willamette River Basin of Western Oregon.
The watershed is situated on the east side of the Oregon Coast Range, southwest of
Corvallis, Oregon (Figure 1).


Figure 1. Location Map (Hulse et al. 1997).

Elevation ranges from approximately 200 feet above sea level to nearly 2,000 feet on the western most edge of the unit. Of the 79,000 acres, eighty percent is privately owned, some seven percent is contained in the Finley National Wildlife refuge, and the remaining thirteen percent's ownership is attributable to the State of Oregon, U.S.D.A. Forest Service, and the Bureau of Land Management. Approximately three percent of the privately owned acreage is zoned ruralresidential, housing some 3,000 residents with typical lot sizes ranging from two to five acres.

There are an estimated 196 mammalian, amphibian, reptilian, and avian species that are thought to currently exist in the watershed. The 196 species modeled represent species that were not locally extirpated or introduced, and preferred at least one of the land-use habitats represented by this study (see Appendix 1 for a complete list of species, diversity weights, and IUCN imperilment ranks). Based on Oregon Natural Heritage (1999) IUCN imperilment rankings, seven of these species are considered imperiled, thirteen are listed as vulnerable, seventy-eight of the species are currently classified as apparently secure, and the remaining ninety-eight species are listed as secure under present conditions. Habitat preferences are variable and range from species that rely on forested and open habitats for nesting and home ranges, to species like the cliff swallow (Hirundo pyrrhonota) that thrive in the commercial/residential communities of the watershed.

Accommodating these wildlife species, the Muddy Creek Watershed encompasses a diverse mix of agriculturally productive, natural-open, residential, and forestry related lands. Elevation, soil type, and slope generally characterize where we
would expect to find such uses, with residential lands scattered amongst the agricultural (higher quality soils) and open (poorer soils) land uses typically observed at lower elevations, and forested lands being observed at higher elevations.

The watershed's northeastern most edge abuts against the Corvallis urban growth boundary, a zone where there has been residential expansion in recent years to accommodate the growing population in the surrounding areas. Transportation routes through the drainage range from a secondary highway, 99 E , to maintained logging roads covering the upland hillsides. Much of the area is zoned for agriculture and forestry related uses, and as can be noted by driving through the area, recent clearcuts and a centrally located mill indicate the community's continued reliance on its timber base as a major source of income.

1990 land-use in the Muddy Creek Watershed is depicted by Figure 2. This illustration comes from the Institute for a Sustainable Environment (ISE), who developed a set of geographic information system (GIS) coverages related to the watershed. These 30 meter grids depict 1990 land-use, 2025 projections of land-use based on trends in use and assumptions regarding future management emphasis, elevation, slope, and soil type.


Figure 2. 1990 Land Use (Hulse et al. 1997).

The ISE's GIS representations of the watershed were further aggregated for modeling purposes. The ISE's classes were further aggregated by the Muddy project habitat team into 26 wildlife habitat classes. Utilizing ArcInfo and ARCView spatial analysis software, six slope-soil based site classes, and four land-use classes; commercial/residential, agriculturally based, open, and forested use were defined.

These site types, and land uses were an aggregate representation of the wildlife
habitat classes. The land use aggregations were the same "use" classes that the Muddy project habitat team adopted to characterize the landscape (see Table 1), and were coincidentally highly correlated with species habitat preferences.

| Wildlife Habitat Classes | Habitat Aggregations |
| :--- | :--- |
| 0-40 Douglas-fir | 0-40 Douglas-fir |
| 40-80 Douglas-fir | 40-80 Douglas-fir |
| 80-120+ Douglas-fir | $80-120+$ Douglas-fir |
| Mixed species forest | Open |
| Deciduous forest | Open |
| Low riparian | N/A |
| Low Marsh | N/A |
| Low stream 1 ${ }^{\text {st }}$ order | N/A |
| Up stream 1 | order |
| Low stream 2 | nd order |
| Up stream 2 | order |
| Low water | N/A |
| Up water | N/A |
| Shrub | N/A |
| Hedge | N/A |
| Oak | Open |
| Prairie | Open |
| Row crops | Open |
| Grass seed production | Agriculture |
| Pasture | Agriculture |
| Christmas tree plantations | Agriculture |
| Hybrid Poplar plantations | Agriculture |
| Orchards | Agriculture |
| Commercial/Residential | Agriculture |
| Roadside | Agriculture |
|  | Residential |
| Site Characteristics | N/A |
| Good soils, slope < 10\% slope |  |
| Good Soils, slope $>10 \%$ slope | Soil-Slope Site Class |
| Moderate soils <10\% slope | 1 |
| Moderate soils $>10 \%$ slope | 2 |
| Poor soils < 10\% slope | 3 |
| Poor soils $>10 \%$ slope | 4 |
|  | 5 |
|  | 6 |

Table 1: Habitat and Soil - Slope Aggregations.

Given these land-use classes, as well as species' preferences for such habitats, the construction of a wildlife diversity index and a supply curve for biodiversity was possible. The methods for achieving this objective follow.

### 2.2 A Marginal Cost Curve for Biodiversity

A marginal cost curve for biodiversity represents a continuum of efficient land allocation opportunities that exist for the Muddy Creek Watershed. This supply curve measures gains in biodiversity against the opportunity cost of foregone land value as lands are reallocated to favor wildlife conservation over other market valued uses. As such, it is representative of meeting a desired level of biological protection at the minimum cost. Utilizing this information a hypothetical land manager would have the ability to make informed, efficient decisions with regards to biodiversity and land allocations. Moreover, this manager would have the ability to choose optimal land allocations for biodiversity given a budget constraint.

This marginal cost curve was revealed by evaluating future land-use allocations and the tradeoffs that existed between biological conservation and commodity production. Biodiversity and market value maximizing solutions were jointly solved by maximizing an additive objective function for total watershed value (see Equation 1). This was accomplished by the development of an objective function capturing both the dynamics of wildlife biodiversity, and market valued uses. Each will be reviewed in turn.

### 2.2.1 Evaluating Biodiversity

Biodiversity by definition reflects a diverse and abundant population of all native species. In modeling multi-species biodiversity it makes intuitive sense to formulate an index that captures both the diversity that an individual species contributes to the overall ecological system, and the probability that the species will actually contribute this diversity to future generations based on current population size and extinction rates.

Montgomery et al. (1999), was able to describe these effects simultaneously: a framework utilized throughout this project to characterize expected wildlife biodiversity. This framework relied on the use of an International Union for Conservation of Nature and Natural Resources (IUCN) Red List Category-based viability function, and the construction of taxonomically based uniqueness rankings.

The viability curve and species uniqueness weights were constructed for 196 of the 234 mammalian, reptilian, avian, and amphibian species, (S), that exist in the Muddy Creek Watershed (Hulse 1997). A general species viability relationship (Vs) was constructed based upon IUCN Red List Category D classes. (These red list categories define degree of imperilment in terms of the expected population size (Ps) of a species and an implied probability of extinction over a defined time horizon (Mace and Lande 1991, Mace and Stuart 1994)). By normalizing these rates of extinction over a single time period, a relationship between population size and probability of survival was determined. A logistic curve was then fitted through the mean population size and the corresponding probability of survival for each of the red list categories, represented by Equation 2. Given the universal nature of this
function, this relationship was used to estimate the viability for each of the species in this study.

$$
\begin{equation*}
\left.\mathrm{V}_{\mathrm{s}}=(1+\exp (-3.2-1.9 * \ln ((\mathrm{Ps}) / 1000)))\right)^{-1} \tag{2}
\end{equation*}
$$

Viability, $\mathrm{V}_{\mathrm{s}}$, within this framework, relies on an implied population size given a certain configuration of habitats or land uses. For example, as more acres of old growth forest are preserved, one might expect the viability of species that prefer such habitat to increase.

To capture such an effect, expected population size, $\mathrm{P}_{\mathrm{s}}$, for each species, s , is dependent on species-specific habitat preferences relative to the Muddy Creek Watershed (White et al. 1997) (see Equation 3). These preferences, determined by a team of biologists familiar with the species of the site, are an integer suitability score of species habitat preference from zero to ten for twenty-six of the land-use categories modeled by the ISE (see Appendix 2 for a complete list habitat preference rankings). Given these habitat preference rankings and implied populations based on the Oregon Nature Conservancy's species specific IUCN endangerment ratings, an average individual per acre measurement, $\mathrm{X}_{\mathrm{sk}}$, was constructed for all species, s , and landuses, k , relative to the study based on current habitat allocations (see Appendix 3). This density index, $\mathrm{X}_{\mathrm{sk}}$, was a fixed measure in the optimization process that required two assumptions for estimation: 1) the proportion of habitat types within each landuse class were fixed, 2) IUCN imperilement rankings were representative of current land allocations in the watershed.
$\mathrm{X}_{\mathrm{sk}}=$ Individual per acre indices for each species in each land-use class.
Where:
$\mathrm{P}_{\mathrm{IUCN}}=$ Original IUCN population estimate
$\mathrm{HP}_{\mathrm{sk}}=$ Habitat Preference Rank

$$
\mathrm{X}_{\mathrm{sk}}=\left[\mathrm{P}_{\mathrm{IUCN}} *\left[\left(\mathrm{HP}_{\mathrm{sk}} * \mathrm{Q}_{\mathrm{k}}\right) /\left(\sum_{\mathrm{k}=1}^{10} \mathrm{HP}_{\mathrm{sk}} * \mathrm{Q}_{\mathrm{k}}\right)\right]^{-1}\right]^{*} \mathrm{Q}_{\mathrm{k}}^{-1}
$$

Population density indices, $\mathrm{P}_{\mathrm{s}}$, were then calculated based on the measure, $\mathrm{X}_{\mathrm{sk}}$, and land-use allotments, $\mathrm{Q}_{\mathrm{jk}}$ (Equation 3).

$$
\begin{equation*}
P_{s}=\sum_{k=1}^{10}\left(X_{s k} * \sum_{j=1}^{6} Q_{j k}\right) \tag{3}
\end{equation*}
$$

## Where:

$P_{s}=$ Species population index
$\mathrm{k}=$ Land-use: residential, agricultural, open, forest
Where: Forest use alternatives include
40 year fully regulated forest
45 year fully regulated forest
50 year fully regulated forest
60 year fully regulated forest
80 year fully regulated forest
120 year fully regulated forest
Park, non-harvested forest
$\mathrm{j}=$ Slope and soil based site class
$\mathrm{Q}_{\mathrm{jk}}=$ Acres of land allocated to each site class and land-use class.

Individual diversity weights, $\mathrm{W}_{\mathrm{s}}$, were constructed for each species, s , based on a hybrid of the taxonomic diversity index proposed by Vane-Wright et al. (1991). A cladogram specific to the Muddy Creek Watershed was first constructed by eliminating all extirpated, and non-watershed related species from an Oregon specific cladogram (Huso 1999).

Initially, the number of species joined at each of the taxonomic tree's nodes were summed. The node counts were then utilized to develop diversity weights (Ws) by taking the inverse of the node count and normalizing to one for the least unique species in each of the four taxonomic divisions of the study: mammalian, reptilian, avian, and amphibian. An example of a cladogram, node counts, and diversity weights for amphibian species present in the Muddy Creek Watershed follows in Figure 3.

For this representation of diversity, species with more close taxonomic relatives have higher node counts, and in turn lower diversity weights. The species with the highest node counts received diversity weights of 1 , the lowest possible rank. In contrast, a unique species like the tailed frog, Ascaphus truei (depicted in Figure 3 ), received a higher diversity weight because of their low relative node count. (The tailed frog's node count of five corresponds to the two branches at the class level, and three divisions at the order level, of this species' taxonomic tree).


Figure 3. Cladogram and Diversity Weights

Expected wildlife biodiversity, $\mathrm{E}(\mathrm{D})$, was then given by the product of individual viability functions, $\mathrm{V}_{\mathrm{s}}\left(\mathrm{P}_{\mathrm{s}}\right)$, and diversity weights, $\mathrm{W}_{\mathrm{s}}$, summed across all species, illustrated by Equation 4.

$$
\begin{equation*}
\mathrm{E}(\mathrm{D})=\sum_{\mathrm{s}=1}^{196} \mathrm{~V}_{\mathrm{s}}\left(\mathrm{P}_{\mathrm{s}}\right) * \mathrm{~W}_{\mathrm{s}} \tag{4}
\end{equation*}
$$

### 2.2.2 Market Values

Market values play an integral role in the development of a marginal cost curve for biodiversity, as these values are used to gauge the opportunity cost of reallocating lands to increase biodiversity protection. But designing a mechanism to accurately reflect the marginal change in watershed land value as an acre(s) of land is converted from one use to another is a difficult exercise.

As a starting point, weighted average land values were determined for the four land-use classes and six soil/slope types. These values were derived by reclassifying and overlaying the ISE's 30 meter grids for 1990 land use, soil type, and slope, with a coverage of tax assessed land values provided to the project by the Benton County Tax Assessors office.

Given these weighted average land values, the opportunity cost of reallocating lands could be directly assigned as the difference in land value between the two uses for which a conversion is made. Several authors have taken this approach including Ando et al. (1998). Relying on average values alone, however, one must assume that marginal land prices are fixed. In other words, even if the entire watershed's land-use was converted to a single use, the marginal value of that land, as well as that of other uses would remain constant and equal to the average value. This assumption is quite limiting, and the only reasonable application would be for situations where management areas are quite small. Thus, converting the entire parcel to a new use would have no impact on how neighboring economies value their land. Presuming that free market forces work, and that a land's currently observed use is its most valued use, this does not seem like a logical assumption to make for this case study.

Instead, a representation of land value that was sensitive to changes in land allocations was needed. The ideal model would accurately reflect changes in the marginal value of land as lands are reallocated, consistent with a classically shaped downward sloping market demand curve for land. O'Sullivan (1993), discusses a useful construct for such an approach, whereby land value functions for each land-use follow a gradient with respect to some variable. O'Sullivan demonstrates this idea utilizing distance from urban center.

For this study, attempts were made to develop these land value functions utilizing elevation as the gradient variable. With the development of such functions, the unobserved portions of the land value curves could be utilized to predict the opportunity cost of reallocating lands. Consider residential lands for example. This land-use is typically observed at lower elevations than forested uses for this watershed. Thus, the marginal value of an additional acre of residential land, is likely going to be much higher at lower elevations where road systems and residential infrastructures are already in place to deal with the needs of such uses. And, an acre of land at the top of a roadless mountain is valued less for residential use, which is unobserved and unknown without estimation, than for its production potential in forestry. Both of these ideas are captured through the development of land value functions.

Hedonic pricing was used to empirically estimate these land value functions. This methodology utilized plot level data on site characteristics and value to estimate land attribute prices (Lopez et al. 1994,Turner et al.1991). For this study, econometric estimation yielded poor results (R-squared(s) of less than .1, and over-
valued residential land-use coefficients). Presuming that we observe an acre's highest valued use, the margins of land area attributable to each land-use did not correspond to this empirical estimation. Ordinary least squares and ordinary least squares corrected for selectivity bias, following the Heckman two-step method, were used to determine land attribute prices. Value was regressed on several combinations of data relating to slope, ownership, zoning, soils, land-use, elevation, and proximity to roads and city centers. Though still an area of continued research, these efforts were ultimately abandoned for this project based on the feasibility of the estimates.

Instead, demand curves were constructed for six slope-soil based site classes and four land-use classes; commercial/residential, agriculturally based, open, and forested use. These linear demand curves, ordered from the highest to lowest average valued use, pass through the weighted average value at the mid quantity of acres currently allocated to each use respectively. The slope of each curve represents the marginal value of an additional acre of land allocated to that use. The exact slope is not known, but was approximated in such a way that it intersected the next highest valued use at the margins of current area (1990) allocation, assuming that a land's observed use is its highest valued use. Each curve's intercept was then shifted to produce the 2025 high development scenario's land allocation across uses (see Appendix 4. For current and 2025 land allocation projections). For forested use this represents high intensity short rotation management. This admittedly crude final set of demand curves reflects the free market forces 2025 outcome for land allocation measured in 1995 dollars (See Figure 4; see Appendix 5 for demand curve coefficients).


Figure 4. Example Demand Curves

### 2.2.3 Forest Valuation

To discern the impact on biodiversity and the associated costs brought about by shifts in forest management, values associated with such changes were estimated. These estimates relied on the ISE's 1990 current land-use coverage which depicted four forested (Douglas-fir, Psuedotsuga menziesii) land-use age classes; 0-40,40 -$80,80-120$, and over 120. Due to the fact that a minimal number of acres was represented by the over 120 years of age class, and also that habitat preference ratings did not discern between the $80-120$ and the greater than 120 year age class, this class was aggregated with the $80-120$ year class for modeling purposes.

These three forested land-use classifications represent all possible land allocation opportunities that exist for modeling biodiversity with respect to forest use. Forest practices were not, however, constrained by rotation ages being equal to 40 , 80, or 120 years. For this analysis, two management intensity classes (Adams 1998) and seven fully regulated forest rotation ages were examined; $40,45,50,60,80,120$,
and park which is a non-harvested forest, see Table 2 . These intensity classes and forest management rotations represent the finest scaled detail this analysis is possible of achieving, given the ISE's original land-use characterizations and species habitat preference ratings. Due to this fact, this project is not able to accurately capture high intensity wildlife habitat management scenarios. Such regimes are targeted at the development of old forest conditions at an earlier age through the use of high intensity thinning strategies. Instead, the broader implications of forest management will be revealed through the steady state analysis of alternative harvest regimes.

|  | Low Intensity Treatment | High Intensity treatment |
| :--- | :---: | :---: |
| Site Prep | X | X |
| Planting density | $400 \mathrm{TPA} ?$ | 432 TPA |
| Pre Commercial thinning <br> Year 13 | $\mathrm{N} / \mathrm{A}$ | Thin to 259 TPA |
| Fertilization year 30 | $\mathrm{N} / \mathrm{A}$ | 200 lbs. Nitrogen |
| Fertilization year 35 | $\mathrm{N} / \mathrm{A}$ | 200 lbs. Nitrogen |
| Commercial thinning <br> Year 35 | $\mathrm{N} / \mathrm{A}$ | Thin to 194 TPA |
| Final harvest | X | X |

Table 2. Management Intensity Classes (Adams 1998).

Each of the forest rotation ages examined represents one of several choices that exist for a landowner when managing forested lands for profit. Private timber companies throughout the Pacific Northwest generally rely on high-intensity, shortrotation forest management. Small private forest holders may or may not choose to mimic the larger corporation's forest practices, and publicly held lands are generally managed for a broader concern for ecological integrity equating to less intensive
management and longer rotations. For each of these lands though, there exists the option for management solely to achieve the objective of profit maximization. Consequently, the ISE chose to appropriate all acres of forested use to a forty-year rotation for their 2025 high development scenario.

To represent the opportunity cost of moving from this "market forces" solution to land allocation strategies contributing more to biodiversity, approximate land and timber values (LTV) for each of these alternative forest management regimes were constructed. First, the soil expectation value (SEV) for each of these seven rotation ages and two management intensity classes was calculated. The SEV measure represents the current bare-ground value of all future costs and returns relating to a specific management regime. Planting, site prep, pre-commercial thinning, and fertilization costs were drawn from Shillinger (1998), log and haul costs were based on TAMM estimates (Adams 2000), and returns were calculated based on expected volumes and Benton County specific log pond prices generated by the Oregon State Forest Service (Corgan 1999), see Tables 3 and 4.

| Associated costs | Low Intensity Treatment | High Intensity treatment |
| :--- | :---: | :---: |
| Site Prep | \$159/acre | \$159/acre |
| Planting | \$119/acre | $\$ 119 / \mathrm{acre}$ |
| Pre Commercial thinning <br> Year 13 | N/A | $\$ 93$ |
| Fertilization year 30 | N/A | $\$ 75$ |
| Fertilization year 35 | N/A | $\$ 75$ |
| Commercial thinning <br> Year 35 | $\mathrm{N} / \mathrm{A}$ | $\$ 168 / \mathrm{MBF}$ |
| Final harvest | \$168/MBF | $\$ 168 / \mathrm{MBF}$ |

Table 3. Forest Management Costs (Shillinger 1998).

| Rotation length | Pond value $\sim$ adjusted for diameter <br> premiums in accordance with age |
| :--- | :---: |
| 40 | $\$ 580 / \mathrm{MBF}$ |
| 45 | $\$ 580 / \mathrm{MBF}$ |
| 50 | $\$ 620 / \mathrm{MBF}$ |
| 60 | $\$ 620 / \mathrm{MBF}$ |
| 80 | $\$ 620 / \mathrm{MBF}$ |
| 120 | $\$ 690 / \mathrm{MBF}$ |

Table 4. Log Pond Values
1998 Log pond values scaled by .9699 , the change in the producer price index (ppi) of lumber and wood products in the U.S. between the years of 1995 and 1998. The figures illustrate the value per MBF (thousand board feet) of Douglas-fir logs minus harvest and transportation costs. The variability between rotation ages represents the instigated quality premiums.

A detailed soil's coverage and King's fifty-year site index map were obtained from the State Soil Geographic Data Base (STATSGO) map from the National Resources Conservation Service (NRCS). This site index map was combined with a reclassed soils and slope coverage to determine soil-slope-site class specific site indices for Douglas-fir forests of the region. These site indices ranged from 112 to 122 and roughly corresponded with Oregon's Forest Inventory and Analysis (FIA) average site index for Benton County forested plots of 121 .

Expected volumes and log diameters associated with each of the management regimes and soil-slope-site classes were ascertained through the use of DFSIM, a growth and yield model developed specifically for Pacific Northwest Douglas-fir forests (Ritchie 1999). Alternative growth and yield models exist, most notably ORGANON, but for this application DFSIM seemed most appropriate based on the feasibility of its projected volumes for rotations longer than 80 years.

Predicted thinning and harvest volumes, as well as $\log$ diameter information, was drawn from DFSIM. Based on these $\log$ diameters and $\log$ pond values, a generalized quality premium was constructed utilizing $\log$ grades characterized by Bell and Dilworth (1988). Based on these quality premiums, harvest volumes, log prices, and costs, SEVs and LTVs were calculated for each management regime. These measures were calculated with an interest rate of seven percent, the average real AAA bond rate over the last decade. The formula used to calculate SEV is depicted by Equation 5, estimated SEVs for each management regime are illustrated by Table 5, and the estimated changes in LTV as forested regimes are shifted from the profit maximizing 40 year rotation to longer rotations are depicted by Table 6 . (Note that these LTV opportunity costs were calculated utilizing an area control approach to account for forest conversions).

$$
\begin{align*}
\mathrm{SEV}=[-\mathrm{PC} & \left.-\frac{\mathrm{PCT}}{(1.07)^{13}}-\frac{\mathrm{FE}}{(1.07)^{30}}-\frac{\left(\mathrm{FE}+\mathrm{LH}^{*} \mathrm{O}\right)}{(1.07)^{35}}+\frac{\mathrm{PQ}^{*} \mathrm{TH}}{(1.07)^{35}}+\frac{\mathrm{PQ} * \mathrm{H}-\mathrm{LH} * \mathrm{Q}}{(1.07)^{\mathrm{t}}}\right] \\
& *\left[\begin{array}{c}
\left.1+\frac{1}{(1.07)^{\mathrm{t}}-1}\right]
\end{array}\right. \tag{5}
\end{align*}
$$

## Where:

SEV = Soil expectation value
PC = Planting and site prep costs
PCT = Pre-commercial thinning cost
FE = Fertilization cost
$\mathrm{PQ}=$ Revenue/MBF timber, scaled with a log diameter quality premium
$\mathrm{TH}=$ Thinning volume harvested (MBF)
LH $=$ Log and Hauling cost/MBF
$\mathrm{H}=$ Final harvest volume (MBF)
$\mathrm{t}=$ Rotation age

| Intensity | Site <br> Index | Site <br> class | $\mathbf{4 0}$ | $\mathbf{4 5}$ | $\mathbf{5 0}$ | $\mathbf{6 0}$ | $\mathbf{8 0}$ | $\mathbf{1 2 0}$ |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Low | 122 | 1 | $\$ 363$ | $\$ 343$ | $\$ 338$ | $\$ 162$ | $-\$ 109$ | $-\$ 252.5$ |
| Low | 122 | 2 | $\$ 363$ | $\$ 343$ | $\$ 338$ | $\$ 162$ | $-\$ 109$ | $-\$ 252.5$ |
| Low | 112 | 3 | $\$ 171$ | $\$ 179$ | $\$ 190$ | $\$ 72$ | $-\$ 137$ | $-\$ 255.9$ |
| Low | 112 | 4 | $\$ 171$ | $\$ 179$ | $\$ 190$ | $\$ 72$ | $-\$ 137$ | $-\$ 255.9$ |
| Low | 119 | 5 | $\$ 315$ | $\$ 300$ | $\$ 298$ | $\$ 137$ | $-\$ 117$ | $-\$ 253.5$ |
| Low | 114 | 6 | $\$ 199$ | $\$ 204$ | $\$ 214$ | $\$ 87$ | $-\$ 132$ | $-\$ 255.2$ |
| High | 122 | 1 | $\$ 605$ | $\$ 588$ | $\$ 575$ | $\$ 368$ | $\$ 48$ | $-\$ 43.4$ |
| High | 122 | 2 | $\$ 605$ | $\$ 588$ | $\$ 575$ | $\$ 368$ | $\$ 48$ | $-\$ 43.4$ |
| High | 112 | 3 | $\$ 372$ | $\$ 378$ | $\$ 376$ | $\$ 201$ | $-\$ 88$ | $-\$ 211.3$ |
| High | 112 | 4 | $\$ 372$ | $\$ 378$ | $\$ 376$ | $\$ 201$ | $-\$ 88$ | $-\$ 211.3$ |
| High | 119 | 5 | $\$ 549$ | $\$ 536$ | $\$ 525$ | $\$ 326$ | $\$ 16$ | $-\$ 81.5$ |
| High | 114 | 6 | $\$ 403$ | $\$ 406$ | $\$ 403$ | $\$ 222$ | $-\$ 72$ | $-\$ 194.4$ |

Table 5. Soil Expectation Values

| Site <br> Index | $\mathbf{4 5}$ | $\mathbf{5 0}$ | $\mathbf{6 0}$ | $\mathbf{8 0}$ | $\mathbf{1 2 0}$ | Park |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 122 | $\$ 294$ | $\$ 299$ | $\$ 470$ | $\$ 1739$ | $\$ 2873$ | $\$ 4248$ |
| 112 | $\$ 201$ | $\$ 148$ | $\$ 226$ | $\$ 1314$ | $\$ 2274$ | $\$ 3505$ |
| 119 | $\$ 272$ | $\$ 267$ | $\$ 419$ | $\$ 1639$ | $\$ 2727$ | $\$ 4065$ |
| 114 | $\$ 213$ | $\$ 166$ | $\$ 255$ | $\$ 1372$ | $\$ 2360$ | $\$ 3618$ |

Table 6. Land and Timber Values

Ultimately, the low intensity management scenario for forest use was dropped, as this management strategy failed to produce higher SEVs or LTVs for any forest rotation length analyzed, than that of any of the high intensity management regimes. Changes in LTV, depicted by Table 6, were then utilized to reduce the market value for lands allocated to alternative forest management regimes. These changes in LTV
measures represented the opportunity (conversion) cost of shifting land from the profit maximizing forty-year rotation, to a longer rotation length.

### 3.0 RESULTS

In developing a marginal cost curve for biodiversity, the opportunity cost of reallocating land from its highest valued use to a new use that has a higher biodiversity value but lower market value is measured against gains in the biodiversity index (Montgomery 1999). This is accomplished by developing an objective function that maximizes land value and the expected biodiversity measure simultaneously (Equation 6).

$$
\begin{equation*}
\mathrm{TV}=\sum_{\mathrm{j}=1}^{6} \sum_{\mathrm{k}=1}^{4}\left[\int_{\mathrm{q}=0}^{\mathrm{Qkj}}\left(\alpha_{\mathrm{kj}}+\beta_{\mathrm{kj}} \mathrm{q}\right) \mathrm{dq}\right]-\sum_{\mathrm{j}=1}^{6} \sum_{\mathrm{m}=1}^{6}\left[\mathrm{C}_{\mathrm{jm}} * \mathrm{Q}_{\mathrm{jm}}\right]+\mathrm{B} *\left(\sum_{\mathrm{s}=1}^{196} \mathrm{~V}_{\mathrm{s}}\left(\mathrm{P}_{\mathrm{s}}\right) * \mathrm{~W}_{\mathrm{s}}\right) \tag{6}
\end{equation*}
$$

Where:
TV = Total value of the Muddy Creek Watershed
$\mathrm{k} \quad=$ Land-use (residential, open, agriculture, seven forest alternatives)
j $\quad=$ Slope and soil-based site class
$\mathrm{m} \quad=$ Forest management regime
$\alpha_{\mathrm{kj}} \quad=$ Demand curve intercept
$\beta_{\mathrm{kj}} \quad=$ Estimated slope of the demand curve
$\mathrm{C}_{\mathrm{kj}}=$ LTV conversion cost
B = Biodiversity scaling weight
$\mathrm{V}_{\mathrm{s}} \quad=$ Species viability
$\mathrm{P}_{\mathrm{s}} \quad=$ Population index (see Equation 3)
$\mathrm{W}_{\mathrm{s}}=$ Diversity weight

Equation 6 represents a total value index of the Muddy Creek Watershed, represented by the sum of the predicted land and biodiversity values. The first expression in the equation denotes the total market value of the watershed, given a particular configuration of uses (k) in site classes (j). The biodiversity value, the second expression in the objective function, is the expected biodiversity measure previously discussed (see Equations 2-4), scaled by (B).

The demand curves were calibrated so that maximizing this objective function without regard to biodiversity (biodiversity weight $(B)=0$ ) yielded the ISE's high development scenario land allocation. This represents the worst-case scenario for biodiversity in the Muddy Creek Watershed. Similarly, dismissing land value from the objective function yields the optimal arrangement of habitats that maximize expected biodiversity. These two scenarios represent the extreme outcomes for the management of biodiversity.

Utilizing the non-linear optimization software package, GAMS MINOS5, intermediate points along the marginal cost curve for biodiversity were identified by solving Equation (6) repeatedly, incrementally increasing the biodiversity weight (B) (See Appendix 6. for an example GAMS code). By varying (B), and simulating landowner supply responses, biodiversity demand was shifted revealing a marginal cost curve for wildlife diversity.

The marginal cost curve for biodiversity, illustrated by figure 5 , depicts all possible wildlife diversity outcomes for this index and future land management opportunities. The expected diversity index ranges from a maximum value of 310.18 to a minimum index value is 296.19 representing the expected biodiversity yielded by the ISE's high development "free market forces" scenario. Current land-use configurations yield a biodiversity index of 307.64. At this level of biodiversity protection, an additional increment in expected diversity would cost landowners nearly ten million dollars. The total cost of moving from no biological protection (high development scenario) to this level of protection is given as the area under the
marginal cost curve, or the integral from a biodiversity index of 296.19 to 307.64 , approximately 22.5 million dollars.

Maintaining the current landscape into 2025 is an inefficient solution to the land allocation problem. To discern this inefficiency, a total opportunity cost curve, representing the accumalated present value of foregone future land rents, was constructed (see Appendix 7). The current landscape yields a biodiversity index of 307.64 , representing a predicted total opportunity cost of 56.8 million dollars. For a similar expenditure on biodiversity protection land managers could, however, come within a fraction of a biodiversity index point of the maximum attainable diversity for this watershed.


Figure 5. Marginal Cost Curve for Biodiversity

In determining the likely impacts of forest management on the biodiversity index, management regimes were tracked throughout the land allocation optimization process. From a modeling standpoint, each acre that is allocated to each of the forest rotations respectively is portioned into one of three habitat associations; 0-40, 40-80, or 80-120 year-old forests. The sixty year rotation for example, assigns two-thirds of its total area to the $0-40$ habitat association, whereas one-third of its area would be contributed to the 40-80 classification. The park scenario allocates one hundred percent of its acres to the 80-129 land-use class.

The resulting forest allocations range from all forested acres being allotted to the forty-year rotation under the free market forces solution, to approximately 4,000 acres allocated to the forty-year management regime and the remaining 17,000 forested acres being deemed park for the maximum attainable biodiversity solution. Intermediate regimes, management rotation lengths of 80 and 120, are not favored over the park scenario for their contribution to the biodiversity index, given an associated gain in LTV (see Figure 6).


Figure 6. Changes in Forest Management

### 4.0 SUPPLEMENTARY ANALYSIS

The marginal cost curve for biodiversity represents all possible efficient land allocation opportunities in the Muddy Creek Watershed when biodiversity is of concern. As such, it is a useful construct to gauge the implications of management strategies targeting imperiled species alone, and the efficiency of land allocations supported by the ISE's possible futures project.

### 4.1 A Marginal Cost Curve for Imperiled Species

The endangerment of native species has been of great national concern over recent decades. Such concern led to adoption of the Endangered Species Act, and has ultimately reshaped federal and state land management planning. Emphasis, and often times priority, is given to those management strategies targeting imperiled species.

A marginal cost curve for imperiled species conveys information relating to the specific needs and preferences of such species. The curve represents land configurations yielding a desired level of imperiled species protection at the minimum cost. By tracking the watershed's total biodiversity as imperiled species diversity is specifically optimized, the implications of such management tactics are revealed.

Seven imperiled species are thought to exist in Muddy Creek Watershed, see Table 7 (Csuti et al. 1997b, Oregon Natural Heritage Program 1998). These species
represent about three and a half percent of the total number of species present in the area, and do not include species that have been introduced or locally extirpated.

| Oregon Nature Conservancy's <br> IUCN endangerment rating | Scientific Name | Common Name |
| :--- | :--- | :--- |
| Imperiled | Branta canadensis | Canada Goose |
| Imperiled | Brachyramphus <br> Marmoratus | Marbled Murrelet |
| Imperiled | Athene cunicularia | Burrowing Owl |
| Imperiled | Ammodramus <br> Savannarum | Grasshopper <br> Sparrow |
| Imperiled | Martes pennanti | Fisher |
| Imperiled | Clemmys maormorata | Western Pond Turtle |
| Imperiled | Chrysemys picta | Painted Turtle |

Table 7. Imperiled Species

In order to compare optimal land management for overall biodiversity protection to optimal management for imperiled species only, Equation 6 was again solved repeatedly for a variety of values for the biodiversity weight (B). For this analysis, the expected diversity measure, $\mathrm{E}(\mathrm{D})$, Equation 4, was replaced with an imperiled species diversity measure, $\mathrm{E}(\mathrm{I})$, Equation 7. This formulation of diversity in effect maximized the expected species richness of all imperiled species, i.

$$
\begin{align*}
& \mathrm{E}(\mathrm{D})=\sum_{\mathrm{s}=1}^{196} \mathrm{~V}_{\mathrm{s}}\left(\mathrm{P}_{\mathrm{s}}\right) * \mathrm{~W}_{\mathrm{s}}  \tag{4}\\
& \mathrm{E}(\mathrm{I})=\sum_{\mathrm{i}=1}^{7} \mathrm{~V}_{\mathrm{i}}\left(\mathrm{P}_{\mathrm{i}}\right) \tag{7}
\end{align*}
$$

As the implied price paid for biodiversity increases to favor imperiled populations, and land is allocated to meet the specific needs of these species, overall biodiversity actually declines steadily. Solutions closest to the free market forces solution, a broader diversity of land uses, yield the highest overall values for expected diversity, while land allocations favoring imperiled species alone devote lands to the agriculture and park uses exclusively. Figure 7 contrasts these gains in imperiled species diversity, $\mathrm{E}(\mathrm{I})$, as lands are allocated to meet the specific needs of such species, with declines in overall expected diversity $\mathrm{E}(\mathrm{D})$ (see Appendix 7 for the marginal cost curve for imperiled species diversity). The population index values for each of the seven imperiled species follows in Figure 8.


Figure 7. Expected Biodiversity vs. Imperiled Diversity.


Figure 8. Imperiled Species Population Indices

### 4.2 Inefficiencies in Land Use Planning

The ISE has been steadily producing and analyzing spatial data-sets of western Oregon. The Muddy Creek Watershed is one of the areas this team of landscape designers has devoted much time to, and is in fact the original source of much of the data utilized by this project. As part of their efforts, a series of possible futures for land-use in the watershed were constructed based on trends in use and emphasis in land management planning goals.

The possible futures project developed five plausible land-use scenarios for the year 2025: current trend, high development, moderate development, high conservation, and moderate conservation. These landscapes were constructed by a team of landscape designers relying on information provided by community representatives, biologists, ecologists, land use planners, and population dynamics modelers. The underlying objective of the project was to discern the implications of
these future landscapes from the perspectives of water quality and wildlife biodiversity; indicators of ecological stability.

The high conservation landscape was constructed by incorporating three primary assumptions about land use into forecasting; limited residential expansion, the introduction of windbreaks, hedgerows, and streamside buffers, and increased rotation length in both hybrid poplar plantations and managed Douglas-fir forests. This predicted landscape represents the ISE's best possible scenario for biodiversity. The fact that the possible futures study did not incorporate specific indicators for biodiversity into their modeling procedures, instead relying on group consensus of "what's best" given all their objectives, suggests that the projected landscape might be an inefficient solution to the land allocation problem. To examine the biological contribution of the "Possible Futures for the Muddy Creek Watershed, Benton County, Oregon" (Hulse et al. 1997) high conservation landscape, and the resulting efficiency of such solutions, the projected landscape's allocation of acres across land uses was utilized to develop measures of expected diversity and total land value relying on the previously discussed solution methodologies. Inherent to this framework, expected population estimates were simultaneously forecasted with expected diversity.

The ISE's high conservation land allocation yielded a biodiversity index value of 308.13. This represents a total opportunity cost or reduction in possible future land rents of $\$ 71.5$ million dollars, and is an inefficient solution to this land allocation problem. The biodiversity index shows marginal gains in wildlife diversity over the current landscape, but as is illustrated by Figure 9, local land managers could come
withina fraction of the maximum attainable biodiversity for a similar expenditure on: biodiversity.


Figure 9. ISE's High Conservation Biodiversity

Five of the seven imperiled apecies actually do better in the current configanation of habitats over the high conservation's land-use allocetivis. The Marbled Murrelet, whose population index more than doubled, and the Fisher, were the two species whose population indices rose given the ISE's high conservation scenario. The impact on less endangered species was variable, and can be viewed in Appendix 8, which depicts population estimates for all 196 species under a variety of land-use treatments.

### 5.0 DISCUSSION

While the wildlife biodiversity index used by Montgomery yields a number of useful results and land management implications for the Muddy Creek Watershed, it is limited in the following ways. The diversity index does not capture species spatial preferences including habitat contiguity, edge effects, core area requirements, or neighboring habitat preference. As such, the index does not prescribe plot-based land prescriptions, and instead suggests larger scale habitat allocations by jointly considering all species' habitat preferences and uniqueness ratings. These habitat allocations suggest the broader management implications when biodiversity is of concern, relying heavily on the biologist's species habitat preference ratings, and the demand curves approximated by this project. These limitations coupled with shortcomings in empirical estimation of land values could help to explain some of the discrepancies between the ISE's high conservation contribution to biodiversity and this model's solution. Further disparities could be in part due to the fact that this study fails to include spatial considerations, likely rendering some results infeasible.

The marginal cost curve for biodiversity depicted by figure 5 , represents the full range of management options for the protection of biodiversity. The shape of this curve, and the underlying land allocations, have a number of interesting management implications relative to the watershed. To retain the current level of biodiversity protection into 2025 , which has an associated biodiversity value of 307.64 , would require a one-time current payment to landowners of 22.5 million dollars.

The marginal cost of an additional unit of biodiversity over the current level of protection is nearly ten million dollars. And capturing the final increment in biodiversity, moving from an expected diversity measure of 309 to 310 , has an associated cost of 400 million dollars. (Payments like these could be viewed as a compensatory reimbursements or necessary tax incentives, policy instruments designed to account for the incurred opportunity costs to landowners of biological protection). This leads to a discussion of the marginal gains associated with increasingly higher amounts paid for biodiversity. For roughly a sixty million dollar expenditure on biodiversity, local land managers could come very close to the maximum attainable biodiversity solution for this watershed. To capture the final increments in the biodiversity index though, managers would have to pay nearly seven times this much, primarily reallocating the acres that are currently used for residential purposes to uses more suitable for biodiversity. In addition, the maximum biodiversity arrangement of land uses brings a fractional change in the population indices of species over other biodiversity favoring scenarios, see Appendix 8, and is a questionable use of resources for local land managers. Montgomery et al. (1999), Montgomery et al. (1994), and Ando et al.(1998) report similar findings.

Land allocations favoring imperiled or critically imperiled species have drastic effects on the overall expected diversity of the region. All seven of these species favor either old forest or agriculturally-based habitats. Land allocations scenarios favoring these species thus focus on either park or agriculture uses, drastically impacting the overall biodiversity index for all species that favor a much broader allocation between uses. Such results indicate the need to consider a more
comprehensive set of species and their related needs when addressing land-use allocations.

The ISE's high conservation landscape results in a number of interesting implications from a biodiversity viewpoint. Recall that the ISE's original charge was to develop a set of feasible land allocations for the year 2025, with the high conservation scenario's emphasis being on improved water quality and biodiversityfriendly land-use allocations. The team considered a broader host of objectives than biodiversity and land value when formulating the high conservation landscape. The depicted future land-use map yields a modest gain in the biodiversity index, from 307.64 to 308.13 , and five of the seven imperiled species actual favor the current configuration of land uses over ISE's suggested landscape.

The final point worth mentioning with regards to this future arrangement of habitats, is the ISE's extension of forest rotations to eighty years and the addition of a number of forest reserve sites to seemingly improve biodiversity values. Data generated based on the biodiversity index of this study indicates that larger gains in overall biodiversity simply are not brought about by increasing forest rotation lengths. In fact, most species favor either young or old forests, indicating that local managers would be better served by allowing some degree of intensive forest management, while trying to procure as many forest reserves as possible. The data does not support the idea that eighty-year rotations are better than forty-year rotations when biodiversity is of concern. It does certainly suggest though, that the no harvest 120 -year-old or older forest is most favored by the populations representative of this study.

Overall, forest management plays an integral role in the protection of biodiversity. Expected wildlife diversity benefits more from no harvest or natural forest reserve areas than from simply lengthening the profit maximizing rotation length beyond 40 years. In fact, based on the species habitat preference ratings, only eight of the 196 species modeled actually prefer the $40-80$ year Douglas-fir land-use allocation to other uses. And, all eight of the species find either the neighboring $0-$ 40 or 80-120 plus Douglas-fir habitat categories nearly as appealing.

### 6.0 CONCLUSION

This project adds to a growing base of literature, which at its core, enhances our understanding of how the extinction rates common today can be lessened in the future through efficient land management strategies. In spite of its limitations, the framework utilized for modeling biodiversity in this project has produced a number of useful results. Key results being: 1) the illustration of the importance of protecting and developing old-growth forest conditions, and 2 ) the demonstration of the pitfalls of considering only imperiled or critically imperiled species in conservation land-use planning.

Incorporating this biodiversity index into spatial analyses is a logical next stage in today's developmental frameworks for modeling wildlife diversity. Currently, the data needed to construct a useful model of this caliber is of limited availability. We are, however, slowly bridging the gap of our understanding so that at some point in the near future, both the multi-species biodiversity studies, like the one presented here, and single species models incorporating a species preference for multiple habitats and the relationships that exists between and amongst these habitats, will be united.

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APPENDICES

## Appendix 1.

IUCN imperilment ranks and diversity weights

|  |  | IUCN Rank | Diversity Weight |
| :---: | :---: | :---: | :---: |
| Ambystoma | gracile | 5 | 1.429 |
| Ambystoma | macrodactylum | 5 | 1.429 |
| Aneides | ferreus | 5 | 1.25 |
| Ensatina | eschscholtzii | 5 | 1.25 |
| Plethodon | dunni | 5 | 1 |
| Plethodon | vehiculum | 5 | 1 |
| Taricha | granulosa | 5 | 1.25 |
| Dicamptodon | tenebrosus | 4 | 1.25 |
| Ascaphus | truei | 4 | 2 |
| Pseudacris | regilla | 5 | 2 |
| Rana | aurora | 4 | 1.25 |
| Ardea | herodias | 5 | 2.684 |
| Butorides | virescens | 5 | 2.684 |
| Branta | canadensis | 2 | 2.684 |
| Aix | sponsa | 5 | 2.684 |
| Anas | platyrhynchos | 5 | 2.318 |
| Lophodytes | cucullatus | 5 | 2.684 |
| Cathartes | aura | 5 | 2.684 |
| Pandion | haliaetus | 5 | 2.684 |
| Elanus | leucurus | 3 | 2.04 |
| Haliaeetus | leucocephalus | 4 | 2.04 |
| Circus | cyaneus | 5 | 2.04 |
| Accipiter | striatus | 5 | 1.821 |
| Accipiter | cooperii | 5 | 1.821 |
| Accipiter | gentilis | 3 | 1.821 |
| Buteo | jamaicensis | 5 | 2.04 |
| Aquila | chrysaetos | 5 | 2.04 |
| Falco | sparverius | 5 | 2.04 |
| Dendragapus | obscurus | 5 | 2.429 |
| Bonasa | umbellus | 5 | 2.429 |
| Oreortyx | pictus | 4 | 2.429 |
| Charadrius | vociferus | 5 | 2.833 |
| Gallinago | gallinago | 5 | 2.55 |
| Brachyramphus | marmoratus | 2 | 2.833 |
| Columba | fasciata | 5 | 2.684 |
| Zenaida | macroura | 5 | 3 |
| Tyto | alba | 5 | 3 |
| Otus | kennicottii | 5 | 2.125 |
| Bubo | virginianus | 5 | 2.125 |
| Glaucidium | gnoma | 4 | 2.125 |
| Athene | cunicularia | 2 | 2.125 |
| Strix | occidentalis | 3 | 1.962 |
| Strix | varia | 5 | 1.962 |
| Asio | otus | 4 | 1.962 |
| Asio | flammeus | 4 | 1.962 |
| Aegolius | acadicus | 4 | 2.125 |
| Chordeiles | minor | 5 | 3.4 |
| Chaetura | vauxi | 5 | 3.4 |

IUCN Rank Diversity Weight

| Calypte | anna | 4 | 3 |
| :---: | :---: | :---: | :---: |
| Selasphorus | rufus | 5 | 3 |
| Melanerpes | lewis | 4 | 2.318 |
| Melanerpes | formicivorus | 3 | 2.318 |
| Sphyrapicus | ruber | 4 | 2.55 |
| Picoides | pubescens | 4 | 2.318 |
| Picoides | villosus | 4 | 2.318 |
| Colaptes | auratus | 5 | 2.55 |
| Dryocopus | pileatus | 4 | 2.55 |
| Contopus | borealis | 4 | 1.244 |
| Contopus | sordidulus | 4 | 1.244 |
| Empidonax | traillii | 4 | 1.214 |
| Empidonax | hammondii | 4 | 1.214 |
| Empidonax | difficilis | 4 | 1.214 |
| Tyrannus | verticalis | 5 | 1.308 |
| Eremophila | alpestris | 5 | 1.417 |
| Progne | subis | 3 | 1.275 |
| Tachycineta | bicolor | 5 | 1.214 |
| Tachycineta | thalassina | 5 | 1.214 |
| Hirundo | pyrrhonota | 5 | 1.214 |
| Hirundo | rustica | 5 | 1.214 |
| Cyanocitta | stelleri | 5 | 1.308 |
| Aphelocoma | californica | 5 | 1.308 |
| Corvus | brachyrhynchos | 5 | 1.244 |
| Corvus | corax | 4 | 1.244 |
| Parus | atricapillus | 5 | 1.342 |
| Parus | rufescens | 5 | 1.342 |
| Psaltriparus | minimus | 5 | 1.417 |
| Sitta | canadensis | 5 | 1.342 |
| Sitta | carolinensis | 4 | 1.342 |
| Certhia | americana | 4 | 1.417 |
| Thryomanes | bewickii | 4 | 1.308 |
| Troglodytes | aedon | 4 | 1.244 |
| Troglodytes | troglodytes | 4 | 1.244 |
| Regulus | satrapa | 4 | 1.417 |
| Sialia | mexicana | 4 | 1.275 |
| Catharus | ustulatus | 5 | 1.275 |
| Turdus | migratorius | 5 | 1.275 |
| Ixoreus | naevius | 4 | 1.275 |
| Chamaea | fasciata | 5 | 1.417 |
| Bombycilla | cedrorum | 5 | 1.417 |
| Vireo | solitarius | 4 | 1.308 |
| Vireo | huttoni | 4 | 1.308 |
| Vireo | gilvus | 5 | 1.308 |
| Vermivora | celata | 5 | 1.214 |
| Dendroica | petechia | 4 | 1.109 |
| Dendroica | coronata | 5 | 1.109 |
| Dendroica | nigrescens | 5 | 1.109 |
| Dendroica | occidentalis | 4 | 1.109 |

IUCN Rank Diversity Weight

| Oporornis | tolmiei | 4 | 1.214 |
| :---: | :---: | :---: | :---: |
| Geothlypis | trichas | 5 | 1.214 |
| Wilsonia | pusilla | 5 | 1.214 |
| Icteria | virens | 4 | 1.214 |
| Piranga | ludoviciana | 5 | 1.417 |
| Pheucticus | melanocephalus | 5 | 1 |
| Passerina | amoena | 4 | 1 |
| Pipilo | maculatus | 4 | 1 |
| Spizella | passerina | 4 | 1 |
| Poecetes | gramineus | 4 | 1 |
| Passerculus | sandwichensis | 5 | 1 |
| Ammodramus | savannarum | 2 | 1 |
| Melospiza | melodia | 5 | 1 |
| Zonotrichia | leucophrys | 5 | 1 |
| Junco | hyemalis | 5 | 1 |
| Agelaius | phoeniceus | 5 | 1 |
| Sturnella | neglecta | 4 | 1 |
| Euphagus | cyanocephalus | 5 | 1 |
| Molothrus | ater | 5 | 1 |
| Icterus | bullockii | 4 | 1 |
| Carpodacus | purpureus | 4 | 1.214 |
| Carpodacus | mexicanus | 5 | 1.214 |
| Loxia | curvirostra | 4 | 1.275 |
| Carduelis | pinus | 5 | 1.186 |
| Carduelis | psaltria | 4 | 1.186 |
| Carduelis | tristis | 4 | 1.186 |
| Coccothraustes | vespertinus | 5 | 1.275 |
| Sorex | vagrans | 4 | 1.714 |
| Sorex | pacificus | 3 | 1.714 |
| Sorex | bendirii | 4 | 1.714 |
| Sorex | trowbridgii | 4 | 1.714 |
| Sorex | sonomae | 5 | 1.714 |
| Neurotrichus | gibbsii | 4 | 2.182 |
| Scapanus | townsendii | 4 | 1.846 |
| Scapanus | orarius | 5 | 1.846 |
| Myotis | lucifugus | 4 | 1.263 |
| Myotis | yumanensis | 3 | 1.263 |
| Myotis | evotis | 3 | 1.263 |
| Myotis | thysanodes | 3 | 1.263 |
| Myotis | volans | 3 | 1.263 |
| Myotis | californicus | 4 | 1.263 |
| Lasionycteris | noctivagans | 4 | 1.846 |
| Eptesicus | fuscus | 4 | 1.846 |
| Lasiurus | cinereus | 4 | 1.846 |
| Plecotus | townsendii | 4 | 1.846 |
| Antrozous | pallidus | 3 | 1.846 |
| Sylvilagus | bachmani | 5 | 2.182 |
| Lepus | americanus | 4 | 2.182 |
| Lepus | californicus | 4 | 2.182 |


| Aplodontia | rufa | IUCN Rank 4 | Diversity Weight $1.714$ |
| :---: | :---: | :---: | :---: |
| Tamias | townsendii | 4 | 1.412 |
| Spermophilus | beecheyi | 5 | 1.412 |
| Sciurus | griseus | 4 | 1.412 |
| Tamiasciurus | douglasii | 5 | 1.5 |
| Glaucomys | sabrinus | 4 | 1.5 |
| Thomomys | mazama | 4 | 1.5 |
| Thomomys | bulbivorus | 5 | 1.5 |
| Castor | canadensis | 4 | 1.714 |
| Peromyscus | maniculatus | 4 | 1.2 |
| Neotoma | fuscipes | 5 | 1.091 |
| Neotoma | cinerea | 5 | 1.091 |
| Clethrionomys | californicus | 4 | 1.2 |
| Phenacomys | albipes | 3 | 1.091 |
| Phenacomys | longicaudus | 4 | 1.091 |
| Microtus | townsendii | 4 | 1 |
| Microtus | longicaudus | 5 | 1 |
| Microtus | oregoni | 4 | 1 |
| Microtus | canicaudus | 5 | 1 |
| Zapus | trinotatus | 4 | 1.714 |
| Erethizon | dorsatum | 5 | 1.714 |
| Canis | latrans | 5 | 1.412 |
| Vulpes | vulpes | 4 | 1.6 |
| Urocyon | cinereoargenteus | 4 | 1.6 |
| Ursus | americanus | 4 | 1.714 |
| Procyon | lotor | 5 | 2 |
| Martes | americana | 3 | 1.263 |
| Martes | pennanti | 2 | 1.263 |
| Mustela | erminea | 5 | 1.2 |
| Mustela | frenata | 5 | 1.2 |
| Spilogale | gracilis | 4 | 1.412 |
| Mephitis | mephitis | 5 | 1.412 |
| Felis | concolor | 4 | 1.5 |
| Lynx | rufus | 4 | 1.5 |
| Cervus | elaphus | 5 | 2.667 |
| Odocoileus | hemionus | 5 | 2.4 |
| Chrysemys | picta | 2 | 3.5 |
| Clemmys | marmorata | 2 | 3.5 |
| Elgaria | coerulea | 5 | 1.556 |
| Elgaria | multicarinata | 5 | 1.556 |
| Sceloporus | occidentalis | 5 | 2 |
| Eumeces | skiltonianus | 5 | 2 |
| Charina | bottae | 4 | 2 |
| Coluber | constrictor | 4 | 1.167 |
| Contia | tenuis | 3 | 1.167 |
| Diadophis | punctatus | 4 | 1.167 |
| Pituophis | catenifer | 5 | 1.167 |
| Thamnophis | ordinoides | 5 | 1 |
| Thamnophis | sirtalis | 5 | 1 |
| Crotalus | viridis | 4 | 2 |

Appendix 2.

Species habitat preference ratings

|  |  | For 0-40 | For 40-80 | For $80+$ | Open | Agr | Res |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Ambystoma | gracile | 2 |  | 10 | 5.83 | 0.00 | 0 |
| Ambystoma | macrodactylum | 3 | 3 | 3 | 3.36 | 3.81 | 3 |
| Aneides | ferreus | 2 | 5 | 8 | 80.86 | 0.00 | 0 |
| Ensatina | eschscholtzii | 6 | 10 | 10 | - 6.87 | 0.23 | 0 |
| Plethodon | dunni | 2 | 10 | 10 | - 2.87 | 0.00 | 0 |
| Plethodon | vehiculum | 3 | 10 | 10 | - 2.01 | 0.00 | 0 |
| Taricha | granulosa | 6 | 9 | 10 | - 8.27 | 1.85 | 0 |
| Dicamptodon | tenebrosus | 2 | 10 | 10 | - 4.08 | - 0.00 | 0 |
| Ascaphus | truei | 0 | 7 | 9 | 9 2.16 | - 0.00 | 0 |
| Pseudacris | regilla | 3 | 4 | 4 | $4 \quad 7.27$ | 3.40 | 4 |
| Rana | aurora | 1 | 4 | 4 | 43.20 | 0.95 | 0 |
| Ardea | herodias | 0 | 3 | 4 | 45.70 | 1.67 | 1 |
| Butorides | virescens | 5 | 8 | 8 | 8.6 .95 | 0.00 | 0 |
| Branta | canadensis | 0 | 0 | 0 | 0.05 | 3.02 | 3 |
| Aix | sponsa | 0 | 0 | 0 | ) 6.74 | 0.00 | 0 |
| Anas | platyrhynchos | 0 | 0 | 0 | 0.00 | 4.59 | 0 |
| Lophodytes | cucullatus | 0 | 1 | 2 | 2.02 | 0.00 | 0 |
| Cathartes | aura | 10 | 4 | 7 | 7.48 | 3.89 | 0 |
| Pandion | haliaetus | 0 | 0 | 2 | 20.00 | 0.00 | 2 |
| Elanus | leucurus | 0 | 0 | 0 | - 1.00 | 2.44 | 0 |
| Haliaeetus | leucocephalus | 0 | 2 | 6 | 60.75 | 0.00 | 0 |
| Circus | cyaneus | 2 | 0 | 0 | - 0.30 | 2.81 | 0 |
| Accipiter | striatus | 4 | 10 | 8 | 8.5 .37 | 0.92 | - 1 |
| Accipiter | cooperii | 3 | 9 | 8 | 87.78 | 0.70 | 3 |
| Accipiter | gentilis | 0 | 4 | 5 | 50.00 | 0.00 | 0 |
| Buteo | jamaicensis | 5 | 3 | 4 | $4 \quad 6.65$ | 3.11 | 2 |
| Aquila | chrysaetos | 2 | 0 | 1 | 0.28 | 0.46 | 0 |
| Falco | sparverius | 3 | 0 | 0 | - 3.74 | 5.15 | 8 |
| Dendragapus | obscurus | 6 | 9 | 10 | ) 2.44 | 0.00 | 0 |
| Bonasa | umbellus | 6 | 7 | 6 | $6 \quad 7.25$ | 0.00 | 0 |
| Oreortyx | pictus | 8 | 0 | 0 | 5.01 | 0.92 | 0 |
| Charadrius | vociferus | 0 | 0 | 0 | 0.00 | 1.01 | 5 |
| Gallinago | gallinago | 0 | 0 | 0 | 0.00 | 0.22 | 0 |
| Brachyramph | marmoratus | 0 | 0 | 1 | 0.00 | 0.00 | 0 |
| Columba | fasciata | 4 | 7 | 6 | - 4.64 | 0.00 | 0 |
| Zenaida | macroura | 8 | 0 | 0 | ) 3.36 | 6.04 | 5 |
| Tyto | alba | 0 | 0 | 1 | 1.04 | 5.04 | 10 |
| Otus | kennicottii | 5 | 8 | 0 | ) 8.59 | 0.00 | 3 |
| Bubo | virginianus | 4 | 8 | 10 | - 8.87 | 3.65 | 3 |
| Glaucidium | gnoma | 3 | 9 | 10 | - 3.46 | 0.00 | 0 |
| Athene | cunicularia | 0 | 0 | 0 | - 0.00 | 0.22 | 0 |
| Strix | occidentalis | 0 | 5 | 10 | 0.00 | 0.00 | 0 |
| Strix | varia | 0 | 5 | 9 | 7.15 | 0.00 | 0 |
| Asio | otus | 4 | 4 | 2 | 2.23 | 0.00 | 0 |
| Asio | flammeus | 0 | 0 | 0 | 0.00 | 0.86 | 0 |

For 0-40 For $40-80$ For $80+$ Open Agr Res

| Aegolius | acadicus | 2 | 9 | 10 | 5.34 | 0.00 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Chordeiles | minor | 6 | 0 | 0 | 0.05 | 0.92 | 0 |
| Chaetura | vauxi | 0 | 0 | 10 | 0.00 | 0.00 | 10 |
| Calypte | anna | 0 | 0 | 0 | 7.03 | 0.00 |  |
| Selasphorus | rufus | 10 | 8 | 7 | 7.57 | 0.00 |  |
| Melanerpes | lewis | 0 | 0 | 0 | 2.62 | 0.00 |  |
| Melanerpes | formicivorus | 0 | 0 | 0 | 3.19 | 0.00 | 0 |
| Sphyrapicus | ruber | 2 | 7 | 7 | 6.52 | 0.00 |  |
| Picoides | pubescens | 2 | 0 | 0 | 8.52 | 0.00 |  |
| Picoides | villosus | 2 | 8 | 10 | 5.51 | 0.00 |  |
| Colaptes | auratus | 3 | 4 | 4 | 5.09 | 1.58 | 9 |
| Dryocopus | pileatus | 2 | 8 | 10 | 3.29 | 0.00 |  |
| Contopus | borealis | 4 | 8 | 10 | 0.86 | 0.00 |  |
| Contopus | sordidulus | 3 | 2 | 2 | 7.93 | 0.00 |  |
| Empidonax | traillii | 8 | 0 | 0 | 0.14 | 0.00 | 0 |
| Empidonax | hammondii | 7 | 10 | 6 | 0.43 | 0.00 | 0 |
| Empidonax | difficilis | 2 | 10 | 10 | 5.89 | 0.00 |  |
| Tyrannus | verticalis | 0 | 0 | 0 | 0.21 | 0.68 | 0 |
| Eremophila | alpestris | 0 | 0 | 0 | 0.00 | 0.69 | 0 |
| Progne | subis | 1 | 0 | 4 | 0.00 | 0.00 | 0 |
| Tachycineta | bicolor | 4 | 2 | 3 | 5.97 | 0.65 | 0 |
| Tachycineta | thalassina | 7 | 0 | 0 | 0.23 | 1.08 | 9 |
| Hirundo | pyrrhonota | 0 | 0 | 0 | 0.00 | 0.00 | 6 |
| Hirundo | rustica | 0 | 0 | 0 | 0.00 | 1.30 | 10 |
| Cyanocitta | stelleri | 8 | 9 | 10 | 3.02 | 0.00 | 0 |
| Aphelocoma | californica | 0 | 0 | 0 | 7.54 | 0.52 | 10 |
| Corvus | brachyrhynchos | 1 | 0 | 0 | 2.22 | 4.50 | 10 |
| Corvus | corax | 2 | 7 | 10 | 2.16 | 0.86 |  |
| Parus | atricapillus | 2 | 1 | 1 | 9.58 | 0.00 | 8 |
| Parus | rufescens | 5 | 10 | 10 | 1.30 | 0.00 | 2 |
| Psaltriparus | minimus | 9 | 0 | 0 | 7.89 | 0.48 |  |
| Sitta | canadensis | 3 | 10 | 10 | 2.47 | 0.00 | 2 |
| Sitta | carolinensis | 0 | 0 | 0 | 6.29 | 0.00 |  |
| Certhia | americana | 1 | 8 | 10 | 5.77 | 0.00 |  |
| Thryomanes | bewickii | 7 | 1 | 0 | 7.64 | 0.23 | 2 |
| Troglodytes | aedon | 7 | 1 | 1 | 6.68 | 0.23 |  |
| Troglodytes | troglodytes | 7 | 9 | 10 | 3.89 | 0.00 | 0 |
| Regulus | satrapa | 3 | 9 | 10 | 3.46 | 0.00 | 0 |
| Sialia | mexicana | 4 | 0 | 0 | 2.27 | 2.03 | 2 |
| Catharus | ustulatus | 8 | 10 | 9 | 4.67 | 0.00 | 0 |
| Turdus | migratorius | 8 | 6 | 5 | 7.74 | 3.91 |  |
| Ixoreus | naevius | 0 | 8 | 10 | 2.59 | 0.00 |  |
| Chamaea | fasciata | 7 | 0 | 0 | 2.13 | 0.00 |  |
| Bombycilla | cedrorum | 3 | 0 | 0 | 4.88 | 0.48 |  |
| Vireo | solitarius | 2 | 3 | 2 | 7.15 | 0.00 |  |

For 0-40 For $40-80$ For $80+$ Open Agr Res

| Vireo | huttoni | 1 | 8 | 6 | 6.80 | 0.00 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Vireo | gilvus | 1 | 0 | 0 | 5.21 | 0.00 | 2 |
| Vermivora | celata | 7 | 0 | 0 | 8.95 | 0.00 | 1 |
| Dendroica | petechia | 0 | 0 | 0 | 0.94 | 0.00 | 0 |
| Dendroica | coronata | 0 | 6 | 7 | 1.73 | 0.00 | 0 |
| Dendroica | nigrescens | 2 | 3 | 2 | 7.29 | 0.00 | 0 |
| Dendroica | occidentalis | 2 | 10 | 10 | 2.59 | 0.00 | 0 |
| Oporornis | tolmiei | 10 | 0 | 0 | 3.62 | 0.00 | 0 |
| Geothlypis | trichas | 0 | 0 | 0 | 1.18 | 1.03 | 0 |
| Wilsonia | pusilla | 9 | 8 | 0 | 4.52 | 0.00 | 0 |
| Icteria | virens | 0 | 0 | 0 | 0.90 | 0.00 | 0 |
| Piranga | ludoviciana | 2 | 8 | 7 | 4.95 | 0.00 | 0 |
| Pheucticus | melanocephal | 1 | 5 | 5 | 7.71 | 0.00 | 0 |
| Pipilo | maculatus | 6 | 1 | 0 | 7.35 | 0.92 | 5 |
| Spizella | passerina | 2 | 0 | 0 | 4.82 | 1.15 | 3 |
| Poecetes | gramineus | 0 | 0 | 0 | 0.37 | 1.34 | 0 |
| Passerculus | sandwichensis | 0 | 0 | 0 | 0.37 | 5.27 | 0 |
| Ammodramu | savannarum | 0 | 0 | 0 | 0.14 | 0.22 | 0 |
| Melospiza | melodia | 7 | 2 | 1 | 5.08 | 2.26 | 6 |
| Zonotrichia | leucophrys | 4 | 0 | 0 | 2.28 | 1.40 | 3 |
| Junco | hyemalis | 6 | 8 | 4 | 4.33 | 1.15 | 2 |
| Agelaius | phoeniceus | 0 | 0 | 0 | 0.48 | 3.94 | 0 |
| Sturnella | neglecta | 0 | 0 | 0 | 0.55 | 2.63 | 0 |
| Euphagus | cyanocephalus | 0 | 0 | 0 | 0.99 | 4.20 | 9 |
| Molothrus | ater | 5 | 3 | 2 | 5.73 | 3.04 | 4 |
| Icterus | bullockii | 1 | 0 | 0 | 2.32 | 0.00 | 2 |
| Carpodacus | purpureus | 6 | 10 | 8 | 6.52 | 0.00 | 0 |
| Carpodacus | mexicanus | 0 | 0 | 0 | 4.63 | 1.41 | 10 |
| Loxia | curvirostra | 1 | 9 | 9 | 2.59 | 0.00 | 0 |
| Carduelis | pinus | 5 | 8 | 9 | 1.30 | 0.00 | 0 |
| Carduelis | psaltria | 0 | 0 | 0 | 4.29 | 1.30 | 2 |
| Carduelis | tristis | 4 | 0 | 0 | 4.47 | 2.37 | 4 |
| Coccothraust | vespertinus | 1 | 6 | 8 | 2.16 | 0.00 | 0 |
| Sorex | vagrans | 4 | 4 | 4 | 6.76 | 3.77 | 2 |
| Sorex | pacificus | 5 | 5 | 5 | 5.84 | 0.00 | 0 |
| Sorex | bendirii | 8 | 8 | 8 | 6.25 | 0.00 | 0 |
| Sorex | trowbridgii | 10 | 10 | 10 | 6.45 | 0.00 | 0 |
| Sorex | sonomae | 5 | 5 | 5 | 5.15 | 0.00 | 0 |
| Neurotrichus | gibbsii | 4 | 6 | 8 | 7.08 | 0.00 | 0 |
| Scapanus | townsendii | 1 | 1 | 1 | 4.48 | 5.92 | 8 |
| Scapanus | orarius | 8 | 8 | 8 | 6.87 | 1.51 | 0 |
| Myotis | lucifugus | 4 | 6 | 8 | 4.65 | 3.23 | 8 |
| Myotis | yumanensis | 2 | 3 | 7 | 3.07 | 1.69 | 4 |
| Myotis | evotis | 6 | 6 | 6 | 4.65 | 1.69 | 2 |

For 0-40 For 40-80 For $80+$ Open Agr Res

| Myotis | thysanodes | 3 | 5 | 7 | 2.79 | 1.23 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Myotis | volans | 1 | 4 | 10 | 2.79 | 1.23 | 0 |
| Myotis | californicus | 3 | 5 | 7 | 3.79 | 1.69 | 1 |
| Lasionycteris | noctivagans | 4 | 3 | 9 | 3.45 | 1.69 | 3 |
| Eptesicus | fuscus | 4 | 3 | 9 | 3.86 | 3.23 | 8 |
| Lasiurus | cinereus | 1 | 3 | 8 | 4.10 | 1.01 | 2 |
| Plecotus | townsendii | 2 | 3 | 3 | 2.09 | 0.77 | 2 |
| Antrozous | pallidus | 0 | 0 | 0 | 1.04 | 0.77 | 2 |
| Sylvilagus | bachmani | 4 | 4 | 4 | 4.99 | 0.92 | 0 |
| Lepus | americanus | 10 | 8 | 6 | 5.69 | 0.00 | 0 |
| Lepus | californicus | 0 | 0 | 0 | 0.74 | 7.54 | 0 |
| Aplodontia | rufa | 10 | 10 | 5 | 3.85 | 0.92 | 0 |
| Tamias | townsendii | 10 | 9 | 10 | 6.79 | 0.00 | 0 |
| Spermophilus | beecheyi | 0 | 0 | 0 | 2.15 | 4.73 | 4 |
| Sciurus | griseus | 0 | 6 | 6 | 6.92 | 0.00 | 2 |
| Tamiasciurus | douglasii | 0 | 9 | 10 | 2.91 | 0.00 | 0 |
| Glaucomys | sabrinus | 0 | 10 | 10 | 4.91 | 0.00 | 5 |
| Thomomys | mazama | 8 | 6 | 0 | 2.28 | 1.61 | 0 |
| Thomomys | bulbivorus | 0 | 0 | 0 | 1.68 | 8.69 | 7 |
| Castor | canadensis | 10 | 10 | 10 | 9.77 | 6.19 | 5 |
| Peromyscus | maniculatus | 9 | 9 | 9 | 9.02 | 3.43 | 3 |
| Neotoma | fuscipes | 0 | 0 | 0 | 7.43 | 0.00 | 0 |
| Neotoma | cinerea | 8 | 8 | 8 | 6.53 | 0.00 | 0 |
| Clethrionomy | californicus | 3 | 8 | 10 | 4.60 | 0.00 | 0 |
| Phenacomys | albipes | 10 | 8 | 8 | 3.46 | 0.00 | 0 |
| Phenacomys | longicaudus | 0 | 5 | 10 | 0.00 | 0.00 | 0 |
| Microtus | townsendii | 0 | 0 | 0 | 1.30 | 1.55 | 0 |
| Microtus | longicaudus | 10 | 0 | 0 | 3.43 | 0.00 | 0 |
| Microtus | oregoni | 10 | 7 | 4 | 4.15 | 0.00 | 0 |
| Microtus | canicaudus | 0 | 0 | 0 | 1.94 | 4.70 | 0 |
| Zapus | trinotatus | 4 | 8 | 10 | 5.97 | 0.00 | 0 |
| Erethizon | dorsatum | 10 | 10 | 10 | 8.34 | 0.79 | 0 |
| Canis | latrans | 8 | 7 | 6 | 7.37 | 5.02 | 5 |
| Vulpes | vulpes | 3 | 2 | 1 | 6.19 | 4.79 | 3 |
| Urocyon | cinereoargenteu | 7 | 8 | 9 | 5.99 | 0.00 | 0 |
| Ursus | americanus | 8 | 8 | 8 | 7.61 | 0.70 | 1 |
| Procyon | lotor | 9 | 9 | 9 | 8.70 | 1.88 | 10 |
| Martes | americana | 7 | 8 | 10 | 4.16 | 0.00 | 0 |
| Martes | pennanti | 5 | 8 | 10 | 1.92 | 0.00 | 0 |
| Mustela | erminea | 8 | 8 | 8 | 7.87 | 1.34 | 0 |
| Mustela | frenata | 8 | 8 | 8 | 7.87 | 1.34 | 0 |
| Spilogale | gracilis | 8 | 6 | 6 | 6.90 | 0.00 | 0 |
| Mephitis | mephitis | 4 | 0 | 2 | 5.15 | 2.56 | 5 |
| Felis | concolor | 10 | 7 | 7 | 8.75 | 5.10 | 2 |
| Lynx | rufus | 8 | 8 | 7 | 8.10 | 2.23 | 2 |


|  |  | For 0-40 For 40-80 For $80+$ | Open | Agr | Res |  |  |
| :--- | :--- | :---: | :--- | :--- | :--- | :--- | :--- |
| Cervus | elaphus | 9 | 6 | 7 | 7.09 | 7.45 | 1 |
| Odocoileus | hemionus | 10 | 7 | 7 | 8.75 | 6.22 | 5 |
| Chrysemys | picta | 0 | 0 | 0 | 0.37 | 1.19 | 0 |
| Clemmys | marmorata | 0 | 0 | 0 | 0.42 | 1.74 | 0 |
| Elgaria | coerulea | 10 | 5 | 5 | 4.21 | 0.70 | 3 |
| Elgaria | multicarinata | 3 | 0 | 0 | 6.14 | 1.35 | 4 |
| Sceloporus | occidentalis | 7 | 2 | 0 | 3.08 | 0.01 | 3 |
| Eumeces | skiltonianus | 2 | 0 | 0 | 2.48 | 0.01 | 3 |
| Charina | bottae | 7 | 3 | 3 | 5.61 | 1.12 | 4 |
| Coluber | constrictor | 3 | 3 | 3 | 5.86 | 3.18 | 6 |
| Contia | tenuis | 4 | 3 | 3 | 7.73 | 1.35 | 4 |
| Diadophis | punctatus | 3 | 3 | 3 | 5.13 | 1.35 | 4 |
| Pituophis | catenifer | 3 | 1 | 1 | 4.19 | 4.17 | 8 |
| Thamnophis | ordinoides | 7 | 5 | 3 | 5.59 | 3.11 | 7 |
| Thamnophis | sirtalis | 4 | 3 | 3 | 4.07 | 3.11 | 7 |
| Crotalus | viridis | 0 | 0 | 0 | 4.86 | 1.35 | 0 |

## Appendix 3.

Individual per acre indices

|  |  | For 0-40 | For 40-80 | For 80+ | Open | Agr | es |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Ambystoma | gracile | 0.0451 | 0.203 | 0.2256 | 0.1315 | 0 | 0 |
| Ambystoma | macrodactylum | 0.0648 | 0.0648 | 0.0648 | 0.1157 | 0.0823 | 0.0648 |
| Aneides | ferreus | 0.0795 | 0.1988 | 0.318 | 0.0343 | 0 | 0 |
| Ensatina | eschscholtzii | 0.1014 | 0.169 | 0.169 | 0.116 | 0.0039 | 0 |
| Plethodon | dunni | 0.0479 | 0.2393 | 0.2393 | 0.0687 | 0 | 0 |
| Plethodon | vehiculum | 0.0701 | 0.2336 | 0.2336 | 0.0469 | 0 | 0 |
| Taricha | granulosa | 0.0888 | 0.1332 | 0.148 | 0.1224 | 0.0274 | 0 |
| Dicamptodon | tenebrosus | 0.0137 | 0.0685 | 0.0685 | - 0.028 | 0 | 0 |
| Ascaphus | truei | 0 | 0.0751 | 0.0966 | 0.0232 | 0 | 0 |
| Pseudacris | regilla | 0.059 | 0.0787 | 0.0787 | 0.143 | 0.0669 | 0.0787 |
| Rana | aurora | 0.0118 | 0.0471 | 0.0471 | 0.0378 | 0.0112 | 0 |
| Ardea | herodias | 0 | 0.1039 | 0.1385 | 0.1975 | 0.058 | 0.0346 |
| Butorides | virescens | 0.1015 | 0.1624 | 0.1624 | 0.1411 | 0 | 0 |
| Branta | canadensis | 0 | 0 | 0 | 0.0001 | 0.0055 | 0.0054 |
| Aix | sponsa | 0 | 0 | 0 | 0.613 | 0 | 0 |
| Anas | platyrhynchos | 0 | 0 | 0 | 0 | 0.1931 | 0 |
| Lophodytes | cucullatus | 0 | 0.0716 | 0.1432 | 0.4308 | 0 | 0 |
| Cathartes | aura | 0.1385 | 0.0554 | 0.0969 | 0.1036 | 0.0539 | 0 |
| Pandion | haliaetus | 0 | 0 | 0.8172 | 0 | 0 | 0.8172 |
| Elanus | leucurus | 0 | 0 | 0 | 0.0087 | 0.0214 | 0 |
| Haliaeetus | leucocephalus | 0 | 0.0528 | 0.1584 | 0.0197 | 0 | 0 |
| Circus | cyaneus | 0.1006 | 0 | 0 | 0.015 | 0.1414 | 0 |
| Accipiter | striatus | 0.0742 | 0.1854 | 0.1483 | 0.0996 | 0.017 | 0.0185 |
| Accipiter | cooperii | 0.057 | 0.171 | 0.152 | - 0.1479 | 0.0134 | 0.057 |
| Accipiter | gentilis | 0 | 0.0361 | 0.0452 | 20 | 0 | 0 |
| Buteo | jamaicensis | 0.0995 | 0.0597 | 0.0796 | 0.1323 | 0.0619 | 0.0398 |
| Aquila | chrysaetos | 0.2328 | 0 | 0.1164 | 40.0323 | 0.0535 | 0 |
| Falco | sparverius | 0.0707 | 0 | 0 | 0.0881 | 0.1214 | 0.1885 |
| Dendragapus | obscurus | 0.1238 | 0.1857 | 0.2063 | 0.0503 | 0 | 0 |
| Bonasa | umbellus | 0.1249 | 0.1458 | 0.1249 | 0.151 | 0 | 0 |
| Oreortyx | pictus | 0.0743 | 0 | 0 | 0.0466 | 0.0085 | 0 |
| Charadrius | vociferus | 0 | 0 | 0 | 0 | 0.15 | 0.7413 |
| Gallinago | gallinago | 0 | 0 | 0 | 0 | 0.1931 | 0 |
| Brachyramphus | marmoratus | 0 | 0 | 0.0325 | - 0 | 0 | 0 |
| Columba | fasciata | 0.1027 | 0.1797 | 0.1541 | 0.1192 | 0.0001 | 0 |
| Zenaida | macroura | 0.1389 | 0 | 0 | 0.0583 | 0.1048 | 0.0868 |
| Tyto | alba | 0 | 0 | 0.0315 | 0.0327 | 0.1589 | 0.3154 |
| Otus | kennicottii | 0.093 | 0.1488 | 0.1674 | 40.1598 | 0 | 0.0558 |
| Bubo | virginianus | 0.0559 | 0.1117 | 0.1397 | 0.1238 | 0.0509 | 0.0419 |
| Glaucidium | gnoma | 0.021 | 0.063 | 0.07 | 0.0242 | 0 | 0 |
| Athene | cunicularia | 0 | 0 | 0 | 0 | 0.0058 | 0 |
| Strix | occidentalis | 0 | 0.0301 | 0.0602 | 0 | 0 | 0 |
| Strix | varia | 0 | 0.1588 | 0.2858 | 0.227 | 0 | 0 |
| Asio | otus | 0.041 | 0.041 | 0.0205 | -0.0536 | 0 | 0 |
| Asio | flammeus | 0 | 0 | 0 | 0 | 0.0579 | 0 |
| Aegolius | acadicus | 0.0138 | 0.062 | 0.0689 | 0.0368 | 0 | 0 |
| Chordeiles | minor | 0.3101 | 0 | 0 | 0.0024 | 0.0475 | 0 |
| Chaetura | vauxi | 0 | 0 | 0.8172 | 0 | 0 | 0.8172 |


|  |  | For 0-40 | or 40-80 | For $80+$ | Open | Agr | Res |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Calypte | anna | 0 | 0 | 0 | 0.1626 | 0 | 0.1155 |
| Selasphorus | rufus | 0.1581 | 0.1265 | 0.1107 | 0.1197 | 0 | 0.0949 |
| Melanerpes | lewis | 0 | 0 | 0 | 0.1839 | 0 | 0 |
| Melanerpes | formicivorus | 0 | 0 | 0 | 0.0765 | 0.0001 | 0 |
| Sphyrapicus | ruber | 0.0153 | 0.0535 | 0.0535 | 0.0498 | 0 | 0.0306 |
| Picoides | pubescens | 0.0301 | 0 | 0 | 0.1282 | 0 | 0.0602 |
| Picoides | villosus | 0.0145 | 0.0578 | 0.0723 | 0.0398 | 0 | 0 |
| Colaptes | auratus | 0.0762 | 0.1016 | 0.1016 | 0.1292 | 0.0402 | 0.2286 |
| Dryocopus | pileatus | 0.0158 | 0.0634 | 0.0792 | 0.0261 | 0 | 0 |
| Contopus | borealis | 0.031 | 0.0619 | 0.0774 | 0.0067 | 0 | 0 |
| Contopus | sordidulus | 0.032 | 0.0213 | 0.0213 | 0.0845 | 0 | 0.0533 |
| Empidonax | traillii | 0.1226 | 0 | 0 | 0.0021 | 0 | 0 |
| Empidonax | hammondii | 0.0454 | 0.0649 | 0.0389 | 0.0028 | 0 | 0 |
| Empidonax | difficilis | 0.0128 | 0.0638 | 0.0638 | 0.0376 | 0 | 0.0064 |
| Tyrannus | verticalis | 0 | 0 | 0 | 0.0539 | 0.1761 | 0 |
| Eremophila | alpestris | 0 | 0 | 0 | 0 | 0.1931 | 0 |
| Progne | subis | 0.0205 | 0 | 0.0818 | 0 | 0 | 0 |
| Tachycineta | bicolor | 0.1327 | 0.0663 | 0.0995 | 0.1981 | 0.0215 | 0 |
| Tachycineta | thalassina | 0.2733 | 0 | 0 | 0.009 | 0.0422 | 0.3514 |
| Hirundo | pyrrhonota | 0 | 0 | 0 | 0 | 0 | 3.319 |
| Hirundo | rustica | 0 | 0 | 0 | 0 | 0.1333 | 1.0285 |
| Cyanocitta | stelleri | 0.1474 | 0.1659 | 0.1843 | 0.0557 | 0 | 0 |
| Aphelocoma | californica | 0 | 0 | 0 | 0.4194 | 0.0286 | 0.556 |
| Corvus | brachyrhynchos | 0.0309 | 0 | 0 | 0.0687 | 0.1391 | 0.3089 |
| Corvus | corax | 0.0157 | 0.055 | 0.0785 | 0.017 | 0.0068 | 0 |
| Parus | atricapillus | 0.0766 | 0.0383 | 0.0383 | 0.367 | 0 | 0.3064 |
| Parus | rufescens | 0.1062 | 0.2124 | 0.2124 | 0.0275 | 0 | 0.0425 |
| Psaltriparus | minimus | 0.2295 | 0 | 0 | 0.2012 | 0.0121 | 0.1785 |
| Sitta | canadensis | 0.0679 | 0.2264 | 0.2264 | 0.056 | 0 | 0.0453 |
| Sitta | carolinensis | 0 | 0 | 0 | 0.1786 | 0 | 0.0284 |
| Certhia | americana | 0.0075 | 0.0603 | 0.0754 | 0.0435 | 0 | 0.0075 |
| Thryomanes | bewickii | 0.0627 | 0.009 | 0 | 0.0684 | 0.0021 | 0.0179 |
| Troglodytes | aedon | 0.0611 | 0.0087 | 0.0087 | 0.0583 | 0.002 | 0.0611 |
| Troglodytes | troglodytes | 0.0394 | 0.0507 | 0.0563 | 0.0219 | 0 | 0 |
| Regulus | satrapa | 0.021 | 0.063 | 0.07 | 0.0242 | 0 | 0 |
| Sialia | mexicana | 0.049 | 0 | 0 | 0.0278 | 0.0248 | 0.0245 |
| Catharus | ustulatus | 0.1372 | 0.1715 | 0.1543 | 0.08 | 0 | 0 |
| Turdus | migratorius | 0.1095 | 0.0821 | 0.0684 | 0.106 | 0.0535 | 0.1095 |
| Ixoreus | naevius | 0 | 0.0752 | 0.0941 | 0.0244 | 0 | 0 |
| Chamaea | fasciata | 0.3429 | 0 | 0 | 0.1044 | 0 | 0 |
| Bombycilla | cedrorum | 0.1536 | 0 | 0 | 0.2499 | 0.0248 | 0.3071 |
| Vireo | solitarius | 0.0237 | 0.0356 | 0.0237 | 0.0849 | 0 | 0 |
| Vireo | huttoni | 0.008 | 0.064 | 0.048 | 0.0544 | 0 | 0 |
| Vireo | gilvus | 0.0868 | 0 | 0 | 0.4521 | 0 | 0.1737 |
| Vermivora | celata | 0.2198 | 0 | 0 | 0.2812 | 0 | 0.0314 |
| Dendroica | petechia | 0 | 0 | 0 | 0.1839 | 0 | 0 |
| Dendroica | coronata | 0 | 0.2597 | 0.3029 | 0.0748 | 0 | 0 |
| Dendroica | nigrescens | 0.0785 | 0.1177 | 0.0785 | 0.2859 | 0 | 0 |
| Dendroica | occidentalis | 0.0145 | 0.0726 | 0.0726 | 0.0188 | 0 | 0 |


|  |  | For 0-40 | For 40-80 | For 80+ | Open | Agr | Res |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Oporornis | tolmiei | 0.0997 | 0 | 0 | 0.0361 | 0 | 0 |
| Geothlypis | trichas | 0 | 0 | 0 | 0.1626 | 0.1419 | 0 |
| Wilsonia | pusilla | 0.1891 | 0.1681 | 0 | 0.0949 | 0 | 0 |
| Icteria | virens | 0 | 0 | 0 | 0.1839 | 0 | 0 |
| Piranga | Iudoviciana | 0.0529 | 0.2116 | 0.1851 | 0.1309 | 0 | 0 |
| Pheucticus | melanocephalus | 0.0321 | 0.1606 | 0.1606 | 0.2476 | 0 | 0 |
| Passerina | amoena | 0 | 0 | 0 | 0.1355 | 0.0152 | 0 |
| Pipilo | maculatus | 0.0513 | 0.0086 | 0 | 0.0628 | 0.0079 | 0.0428 |
| Spizella | passerina | 0.0307 | 0 | 0 | 0.074 | 0.0176 | 0.046 |
| Poecetes | gramineus | 0 | 0 | 0 | 0.0147 | 0.0533 | 0 |
| Passerculus | sandwichensis | 0 | 0 | 0 | 0.0132 | 0.189 | 0 |
| Ammodramus | savannarum | 0 | 0 | 0 | 0.0031 | 0.0048 | 0 |
| Melospiza | melodia | 0.1581 | 0.0452 | 0.0226 | 0.1148 | 0.0511 | 0.1355 |
| Zonotrichia | leucophrys | 0.1857 | 0 | 0 | 0.1058 | 0.0649 | 0.1393 |
| Junco | hyemalis | 0.1194 | 0.1592 | 0.0796 | 0.0861 | 0.0229 | 0.0398 |
| Agelaius | phoeniceus | 0 | 0 | 0 | 0.0226 | 0.186 | 0 |
| Sturnella | neglecta | 0 | 0 | 0 | 0.0114 | 0.0544 | 0 |
| Euphagus | cyanocephalus | 0 | 0 | 0 | 0.0378 | 0.1611 | 0.3453 |
| Molothrus | ater | 0.1061 | 0.0637 | 0.0424 | 0.1216 | 0.0644 | 0.0849 |
| Icterus | bullockii | 0.044 | 0 | 0 | 0.1023 | 0 | 0.0881 |
| Carpodacus | purpureus | 0.0324 | 0.054 | 0.0432 | 0.0352 | 0 | 0 |
| Carpodacus | mexicanus | 0 | 0 | 0 | 0.2593 | 0.0789 | 0.5599 |
| Loxia | curvirostra | 0.0084 | 0.0756 | 0.0756 | 0.0218 | 0 | 0 |
| Carduelis | pinus | 0.1221 | 0.1953 | 0.2198 | 0.0316 | 0 | 0 |
| Carduelis | psaltria | 0 | 0 | 0 | 0.0899 | 0.0272 | 0.0419 |
| Carduelis | tristis | 0.0394 | 0 | 0 | 0.044 | 0.0234 | 0.0394 |
| Coccothraustes | vespertinus | 0.0368 | 0.221 | 0.2946 | 0.0795 | 0 | 0 |
| Sorex | vagrans | 0.0223 | 0.0223 | 0.0223 | 0.0377 | 0.0211 | 0.0112 |
| Sorex | pacificus | 0.0165 | 0.0165 | 0.0165 | 0.0193 | 0 | 0 |
| Sorex | bendirii | 0.0433 | 0.0433 | 0.0433 | 0.0339 | 0 | 0 |
| Sorex | trowbridgii | 0.0448 | 0.0448 | 0.0448 | 0.0289 | 0 | 0 |
| Sorex | sonomae | 0.1365 | 0.1365 | 0.1365 | 0.1405 | 0 | 0 |
| Neurotrichus | gibbsii | 0.0283 | 0.0424 | 0.0565 | 0.0501 | 0 | 0 |
| Scapanus | townsendii | 0.0065 | 0.0065 | 0.0065 | 0.0292 | 0.0386 | 0.0522 |
| Scapanus | orarius | 0.1246 | 0.1246 | 0.1246 | 0.107 | 0.0235 | 0 |
| Myotis | lucifugus | 0.0208 | 0.0312 | 0.0417 | 0.0242 | 0.0168 | 0.0417 |
| Myotis | yumanensis | 0.0075 | 0.0113 | 0.0264 | 0.0116 | 0.0064 | 0.0151 |
| Myotis | evotis | 0.0148 | 0.0148 | 0.0148 | 0.0114 | 0.0042 | 0.0049 |
| Myotis | thysanodes | 0.0104 | 0.0173 | 0.0242 | 0.0097 | 0.0043 | 0 |
| Myotis | volans | 0.0039 | 0.0157 | 0.0393 | 0.011 | 0.0048 | 0 |
| Myotis | californicus | 0.0222 | 0.0371 | 0.0519 | 0.0281 | 0.0125 | 0.0074 |
| Lasionycteris | noctivagans | 0.0299 | 0.0224 | 0.0672 | 0.0257 | 0.0126 | 0.0224 |
| Eptesicus | fuscus | 0.0238 | 0.0179 | 0.0536 | 0.023 | 0.0193 | 0.0477 |
| Lasiurus | cinereus | 0.0103 | 0.0308 | 0.0822 | 0.0421 | 0.0103 | 0.0205 |
| Plecotus | townsendii | 0.0266 | 0.04 | 0.04 | 0.0278 | 0.0103 | 0.0266 |
| Antrozous | pallidus | 0 | 0 | 0 | 0.0207 | 0.0153 | 0.0397 |
| Sylvilagus | bachmani | 0.1127 | 0.1127 | 0.1127 | 0.1407 | 0.0259 | 0 |
| Lepus | americanus | 0.0522 | 0.0418 | 0.0313 | 0.0297 | 0 | 0 |
| Lepus | californicus | 0 | 0 | 0 | 0.0055 | 0.0562 | 0 |


|  |  | For 0-40 | For 40-80 | For $80+$ | Open | Agr | Res |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Aplodontia | rufa | 0.0477 | 0.0477 | 0.0238 | 0.0184 | 0.0044 | 0 |
| Tamias | townsendii | 0.046 | 0.0414 | 0.046 | 0.0312 | 0 | 0 |
| Spermophilus | beecheyi | 0 | 0 | 0 | 0.0735 | 0.162 | 0.1371 |
| Sciurus | griseus | 0 | 0.0575 | 0.0575 | 0.0664 | 0 | 0.0192 |
| Tamiasciurus | douglasii | 0 | 0.2593 | 0.2881 | 0.0837 | 0 | 0 |
| Glaucomys | sabrinus | 0 | 0.0718 | 0.0718 | 0.0352 | 0 | 0.0359 |
| Thomomys | mazama | 0.1768 | 0.1326 | 0 | 0.0505 | 0.0355 | 0 |
| Thomomys | bulbivorus | 0 | 0 | 0 | 0.0101 | 0.0523 | 0.0421 |
| Castor | canadensis | 0.0944 | 0.0944 | 0.0944 | 0.0922 | 0.0584 | 0.0472 |
| Peromyscus | maniculatus | 0.1069 | 0.1069 | 0.1069 | 0.1072 | 0.0407 | 0.0356 |
| Neotoma | fuscipes | 0 | 0 | 0 | 0.1839 | 0 | 0 |
| Neotoma | cinerea | 0.1433 | 0.1433 | 0.1433 | 0.1169 | 0 | 0 |
| Clethrionomys | californicus | 0.0212 | 0.0566 | 0.0707 | 0.0325 | 0 |  |
| Phenacomys | albipes | 0.0225 | 0.018 | 0.018 | 0.0078 | 0 |  |
| Phenacomys | longicaudus | 0 | 0.0723 | 0.1446 | 0 | 0 |  |
| Microtus | townsendii | 0 | 0 | 0 | 0.0383 | 0.0459 |  |
| Microtus | longicaudus | 0.3358 | 0 | 0 | 0.115 | 0 |  |
| Microtus | oregoni | 0.0591 | 0.0413 | 0.0236 | 0.0245 | 0 |  |
| Microtus | canicaudus | 0 | 0 | 0 | 0.0704 | 0.1709 |  |
| Zapus | trinotatus | 0.0255 | 0.051 | 0.0637 | 0.038 | 0 |  |
| Erethizon | dorsatum | 0.1348 | 0.1348 | 0.1348 | 0.1125 | 0.0107 | 0 |
| Canis | latrans | 0.0993 | 0.0869 | 0.0745 | 0.0915 | 0.0623 | 0.0621 |
| Vulpes | vulpes | 0.0185 | 0.0123 | 0.0062 | 0.0382 | 0.0296 | 0.0185 |
| Urocyon | cinereoargenteus | 0.0393 | 0.0449 | 0.0505 | 0.0336 | 0 | 0 |
| Ursus | americanus | 0.039 | 0.039 | 0.039 | 0.0371 | 0.0034 | 0.0049 |
| Procyon | lotor | 0.1157 | 0.1157 | 0.1157 | 0.1119 | 0.0242 | 0.1286 |
| Martes | americana | 0.017 | 0.0195 | 0.0243 | 0.0101 | 0 | 0 |
| Martes | pennanti | 0.0035 | 0.0056 | 0.007 | 0.0013 | 0 | 0 |
| Mustela | erminea | 0.1232 | 0.1232 | 0.1232 | 0.1211 | 0.0206 | 0 |
| Mustela | frenata | 0.1232 | 0.1232 | 0.1232 | 0.1211 | 0.0206 | 0 |
| Spilogale | gracilis | 0.048 | 0.036 | 0.036 | 0.0414 | 0 | 0 |
| Mephitis | mephitis | 0.1154 | 0 | 0.0577 | 0.1486 | 0.0737 | 0.1442 |
| Felis | concolor | 0.0341 | 0.0238 | 0.0238 | 0.0298 | 0.0174 | 0.0068 |
| Lynx | rufus | 0.0345 | 0.0345 | 0.0302 | 0.0349 | 0.0096 | 0.0086 |
| Cervus | elaphus | 0.0973 | 0.0649 | 0.0757 | 0.0767 | 0.0805 | 0.0108 |
| Odocoileus | hemionus | 0.1055 | 0.0738 | 0.0738 | 0.0923 | 0.0656 | 0.0527 |
| Chrysemys | picta | 0 | 0 | 0 | 0.0016 | 0.0053 | 0 |
| Clemmys | marmorata | 0 | 0 | 0 | 0.0013 | 0.0054 | 0 |
| Elgaria | coerulea | 0.1933 | 0.0966 | 0.0966 | 0.0813 | 0.0136 | 0.058 |
| Elgaria | multicarinata | 0.1177 | 0 | 0 | 0.2411 | 0.053 | 0.157 |
| Sceloporus | occidentalis | 0.2543 | 0.0727 | 0 | 0.112 | 0.0003 | 0.109 |
| Eumeces | skiltonianus | 0.2032 | 0 | 0 | 0.2522 | 0.001 | 0.3048 |
| Charina | bottae | 0.0491 | 0.021 | 0.021 | 0.0394 | 0.0078 | 0.0281 |
| Coluber | constrictor | 0.0201 | 0.0201 | 0.0201 | 0.0392 | 0.0213 | 0.0402 |
| Contia | tenuis | 0.0124 | 0.0093 | 0.0093 | 0.0241 | 0.0042 | 0.0124 |
| Diadophis | punctatus | 0.0269 | 0.0269 | 0.0269 | 0.0459 | 0.0121 | 0.0358 |
| Pituophis | catenifer | 0.0726 | 0.0242 | 0.0242 | 0.1014 | 0.1009 | 0.1935 |
| Thamnophis | ordinoides | 0.1195 | 0.0854 | 0.0512 | 0.0955 | 0.0531 | 0.1195 |
| Thamnophis | sirtalis | 0.0905 | 0.0679 | 0.0679 | 0.092 | 0.0703 | 0.1584 |
| Crotalus | viridis |  |  |  | 0.0977 | 0.0272 |  |

## Appendix 4.

Current and high conservation land allocations

## Original acres 1990

|  | For 0-40 | For 40-80 For 80+ |  |  |  |  | Open |  | Agr | Res |  |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | :---: | :---: | :---: | :---: |
| s1ss1 | 1085.7 | 1067.5 | 98.7 | 1334.8 | 10638.7 | 642.7 |  |  |  |  |  |
| s1ss2 | 2057.1 | 1987.8 | 410.3 | 1264.3 | 1742.7 | 224.6 |  |  |  |  |  |
| s2ss1 | 673.0 | 685.9 | 233.7 | 1214.5 | 10787.4 | 283.3 |  |  |  |  |  |
| s2ss2 | 3628.1 | 3137.8 | 1667.1 | 1663.5 | 585.1 | 94.1 |  |  |  |  |  |
| s3ss1 | 672.1 | 766.8 | 143.0 | 477.7 | 1112.0 | 110.1 |  |  |  |  |  |
| s3ss2 | 3981.1 | 3880.3 | 2059.1 | 2202.4 | 1024.1 | 151.7 |  |  |  |  |  |

Where:

S1ss1 $=$ good soils, $<10 \%$ slope
S1ss2 $=$ good soils, $>10 \%$ slope
S2ss1 = moderate soils, $<10 \%$ slope
S2ss2 $=$ moderate soils, $>10 \%$ slope
S3ss1 $=$ poor soils, $<10 \%$ slope
S3ss2 $=$ poor soils, $>10 \%$ slope

## 2025 ISE High Conservation Land Allocations:



Courtesy: Hulse et al. 1997

## Appendix 5.

Demand curve coefficients

Original Adjusted for High Development

| s1ss1 | alpha | beta | alpha | beta |
| :---: | :---: | :---: | :---: | :---: |
| res | 7227 | -7.0589 | 9573 | -7.0589 |
| open | 3150.5 | -0.7162 | 3283.2 | -0.7162 |
| ag | 1914.7 | -0.0912 | 1917.6 | -0.0912 |
| for | 1686.4 | -0.0731 | 1686.4 | -0.0731 |
| s1ss2 |  |  |  |  |
| res | 4934 | -15.448 | 8122.2 | -15.448 |
| ag | 1479.5 | -0.0673 | 1485.4 | -0.0673 |
| open | 1420.3 | -0.0372 | 1421.9 | -0.0372 |
| for | 1371.8 | -0.0222 | 1371.8 | -0.0222 |
| s2ss1 |  |  |  |  |
| res | 4728.4 | -11.717 | 7047.9 | -11.717 |
| open | 1469.5 | -0.2141 | 1426.4 | -0.2141 |
| ag | 1207.6 | -0.0393 | 1207.6 | -0.0393 |
| for | 853.17 | -0.0104 | 588.6 | -0.0104 |
| s2ss2 |  |  |  |  |
| res | 3649.9 | -11.611 | 4633.3 | -11.611 |
| ag | 2675.1 | -1.2516 | 2726.2 | -1.2516 |
| open | 2387.5 | -0.8281 | 2408.9 | -0.8281 |
| for | 474.72 | -0.0116 | 474.2 | -0.0116 |
| s3ss1 |  |  |  |  |
| res | 5806 | -32.553 | 11524.1 | -32.553 |
| ag | 2295.9 | -0.6726 | 2144.9 | -0.6726 |
| open | 1844.8 | -0.4188 | 1779.4 | -0.4188 |
| for | 1470.8 | -0.253 | 1470.8 | -0.253 |
| s3ss2 |  |  |  |  |
| res | 3721 | -11.471 | 6896.6 | -11.471 |
| ag | 2059.2 | -0.5165 | 2101.8 | -0.5165 |
| open | 2009.5 | -0.4741 | 2044.2 | -0.4741 |
| for | 444.51 | -0.0109 | 444.5 | -0.0109 |

Appendix 6.

GAMS MINOS5 code

## \$TITLE MUDDY CREEK

## \$OFFUPPER

\$OFFSYMXREF

SET HAB HABITAT
/ncon, mcon, ocon, open, agr, res/;
set hab2 new habs with seven for choices
/ f40, f45, f50, f60, f80, f120, park, openl, agrl, resl/;

SET ELSO SITE PRODUCTIVITY SOIL SLOPE
/S1SS1, S1SS2, S2SS1, S2SS2, S3SS1, S3SS2
/;

## SET S SPECIES

/ al,
a2,
a3,
a4,
a196
/;

TABLE AC(ELSO,HAB2) ACRES IN EACH HABITAT CLASS \$INCLUDE "k:\shunkn\GAMS $\backslash$ ACRES.TXT";
table acl(elso,hab) acres original in each ncon mcon etc \$include " k :\shunkn\gamslacresl.txt";
table qmidl(elso,hab2) original acres mid points
Sinclude "k:\shunkn\gamsimidl.txt";
TABLE alpha(ELSO,HAB2) land value alphas FOR EACH HABITAT CLASS \$INCLUDE "k:\shunkn\GAMS $\backslash a l p h a . T X T " ;$

TABLE beta(ELSO,HAB2) land value betas ag open res equal 0
\$INCLUDE "k:\shunkn\GAMS\beta.TXT";

TABLE adjust(ELSO,HAB2) percent of lv sev adjustments ag open res equal 1 \$INCLUDE "k:\shunkn\GAMSไadjust.TXT";

TABLE SPAC(S,HAB) SPECIES PER ACRE
\$INCLUDE "k:\shunkn\GAMS\SPACl1.TXT";
PARAMETERS TOT(ELSO) TOTAL AREA
/ S1SS1 14868.15
S1SS2 7686.83
S2SS1 13877.83
S2SS2 10775.66
S3SS1 3281.65
S3SS2 13298.72
/;

## PARAMETERS WT(S) SPECIE WEIGHTS

| al | 1.429 |
| :---: | :---: |
| a2 | 1.429 |
| a3 | 1.25 |
| a196 | $2 /$ |

SCALAR BIOPRICE / 1000/;
SCALAR MPRICE / 1/;
SCALAR ALPH /3.2/;
SCALAR BET / 1.9 /;
SCALAR THETA / 1000/;
VARIABLES
ADAC2(ELSO,HAB2) ADJUSTED ACRES
MBVAL;
POSITIVE VARIABLE ADAC2;
EQUATIONS
EQ1(ELSO) CANT MOVE TO NEW ELSO
OBJ;
EQ1(ELSO)..
$\operatorname{SUM}(\mathrm{HAB} 2, \mathrm{ADAC2}(E L S O, H A B 2))=1=\mathrm{TOT}(E L S O)$;

OBJ..
MBVAL $=\mathrm{E}=\mathrm{BIOPRICE} * \operatorname{SUM}(\mathrm{~S}, \mathrm{WT}(\mathrm{S}) /(1+\mathrm{EXP}(-\mathrm{ALPH}-\mathrm{BET} *$
LOG(. $01+(1 /$ THETA $) ~ * ~$
(
(spac(s,"res")* sum(elso,adac2(elso,"res1"))) +
$(\operatorname{spac}(s, "$ open" $) * \operatorname{sum}($ elso, adac2(elso,"open1"))) +
$($ spac(s,"agr") * sum(elso, adac2(elso,"agr1")) ) +
(spac(s,"ncon") * sum(elso,((adac2(elso,"f40") + .888*adac2(elso,"f45") + .8*adac2(elso,"f50") + $.6666^{*}$ adac2(elso,"f60") + . 5 * adac2(elso,"f80") + . $3333^{*}$ adac2(elso,"f120")))) +
(spac(s,"mcon") * sum(elso,((.5*adac2(elso,"f80") + .111* adac2(elso,"f45") + .2 * adac2(elso,"f50") + .333* adac2(elso,"f60") + .333 * adac2(elso,"f120")))) +
$(\operatorname{spac}(s, "$ ocon") * sum(elso, ((.333 * adac2(elso,"f120") + adac2(elso,"park"))))
) )) ) ) ) ) )

+ MPRICE *
(
(sum(elso, (adac2(elso,"res1")/2 * beta(elso,"res1") + alpha(elso,"res1") * adjust(elso,"res1")) *

```
adac2(elso,"res1"))) +
(adac2("s1ss1","open1") * ((adac2("s1ss1","res1") + adac2("s1ss1","open1")/2) *
beta("s1ss1","open1") + alpha("s1ss1","open1")*adjust("s1ss1","open1"))) +
(adac2("s1ss2","agr1") * ((adac2("s1ss2","res1") + adac2("s1ss2","agr1")/2) *
beta("s1ss2","agr1") + alpha("s1ss2","agrl")*adjust("s1ss2","agr1"))) +
(adac2("s2ss1","open1") * ((adac2("s2ss1","res1") + adac2("s2ss1","open1")/2) *
beta("s2ss1","open1") + alpha("s2ss1","open1")* adjust("s2ss1","open1"))) +
(adac2("s2ss2","agr1") * ((adac2("s2ss2","res1") + adac2("s2ss2","agr1")/2) *
beta("s2ss2","agr1") + alpha("s2ss2","agr1") * adjust("s2ss2","agr1"))) +
(adac2("s3ss1","agr1") * ((adac2("s3ss1","res1") + adac2("s3ss1","agr1")/2) *
beta("s3ss1","agr1") + alpha("s3ss1","agr1")* adjust("s3ss1","agr1"))) +
(adac2("s3ss2","agr1") * ((adac2("s3ss2","res1") + adac2("s3ss2","agr1")/2) *
beta("s3ss2","agr1") + alpha("s3ss2","agr1") * adjust("s3ss2","agr1"))) +
(adac2("s1ss1","agr1") * ((adac2("s1ss1","res1") + adac2("s1ss1","open1") +
adac2("slss1","agrl")/2) * beta("s1ss1","agr1") + alpha("s1ss1","agr1") *
adjust("slss1","agrl"))) +
(adac2("s1ss2","open1") * ((adac2("s1ss2","res1") + adac2("s1ss2","agr1") +
adac2("slss2","open1")/2) * beta("s1ss2","open1") + alpha("s1ss2","open1") *
adjust("s1ss2","open1"))) +
(adac2("s2ss1","agr1") * ((adac2("s2ss1","res1") + adac2("s2ss1","openl") +
adac2("s2ss1","agrl")/2) * beta("s2ss1","agrl") + alpha("s2ss1","agrl") *
adjust("s2ssl","agrl"))) +
(adac2("s2ss2","open1") * ((adac2("s2ss2","res1") + adac2("s2ss2","agrl") +
adac2("s2ss2","open1")/2) * beta("s2ss2","open1") + alpha("s2ss2","open1") *
adjust("s2ss2","open1"))) +
(adac2("s3ss1","open1") * ((adac2("s3ss1","res1") + adac2("s3ss1","agrl") +
adac2("s3ss1","open1")/2) * beta("s3ss1","open1") + alpha("s3ss1","open1") *
adjust("s3ss1","open1"))) +
(adac2("s3ss2","open1") * ((adac2("s3ss2","res1") + adac2("s3ss2","agrl") +
adac2("s3ss2","open1")/2) * beta("s3ss2","open1") + alpha("s3ss2","open1") *
adjust("s3ss1","openl"))) +
```

sum(elso,(adac2(elso,"f40") * (((adac2(elso,"res1") + adac2(elso,"open1") + adac2(elso,"agr1") +
(adac2(elso,"f40")+
adac2(elso,"f45")+ adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"f120") +
adac2(elso,"park"))/2) * beta(elso,"f40") + alpha(elso,"f40") * adjust(elso,"f40")) )) ) +

```
sum(elso,(adac2(elso,"f45") * (((adac2(elso,"res1") + adac2(elso,"openl") + adac2(elso,"agr1") +
(adac2(elso,"f40")+
adac2(elso,"f45")+ adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"f120") +
adac2(elso,"park"))/2) * beta(elso,"f45") + alpha(elso,"f45") * adjust(elso,"f45"))))) +
```

sum(elso,(adac2(elso,"f50") * (((adac2(elso,"res1") + adac2(elso,"openl") + adac2(elso,"agrl") + (adac2(elso,"f40")+ adac2(elso,"f45")+ adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"f120") + adac2(elso,"park"))/2) * beta(elso,"f50") + alpha(elso,"f50") * adjust(elso,"f50"))))) +

```
sum(elso,(adac2(elso,"f60") * (((adac2(elso,"res1") + adac2(elso,"openl") + adac2(elso,"agrl") +
(adac2(elso,"f40")+
adac2(elso,"f45") + adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"f120") +
adac2(elso,"park"))/2) * beta(elso,"f60") + alpha(elso,"f60") * adjust(elso,"f60"))))) +
```

```
sum(elso,(adac2(elso,"f80") * (((adac2(elso,"resl") + adac2(elso,"openl") + adac2(elso,"agrl") +
(adac2(elso,"f40")+
adac2(elso,"f45")+ adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"fl20") +
adac2(elso,"park"))/2) * beta(elso,"f80") + alpha(elso,"f80") * adjust(elso,"f80"))))) +
sum(elso,(adac2(elso,"f120") * (((adac2(elso,"resl") + adac2(elso,"open1") + adac2(elso,"agrl") +
(adac2(elso,"f40")+
adac2(elso,"f45")+ adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"f120") +
adac2(elso,"park"))/2) * beta(elso,"f120") + alpha(elso,"f120") * adjust(elso,"f120"))))) +
sum(elso,(adac2(elso,"park") * (((adac2(elso,"resl") + adac2(elso,"openl") + adac2(elso,"agrl") +
(adac2(elso,"f40")+
adac2(elso,"f45")+ adac2(elso,"f50") + adac2(elso,"f60") + adac2(elso,"f80") + adac2(elso,"f120") +
adac2(elso,"park"))/2) * beta(elso,"park") + alpha(elso,"park") * adjust(elso,"park")))))
)
- sum(elso, sum(hab2, (adac2(elso, hab2) * adjust(elso, hab2))));
```

MODEL MUDDY / ALL /;
SOLVE MUDDY USING NLP MAXIMIZING MBVAL;
PARAMETERS
olv original land value
NLV NEW LAND VALUE
OBIO ORIGINAL BIO VALUE
NBIO NEW BIO VALUE
EXPD EXPECTED DIVERSITY
AB(S) SPECIES POPULATION INDEX
NAB(S) NEW SPECIES POPULATION INDEX
AREA TOTAL ACRES;

```
olv = (sum(elso, (ac(elso,"resl")/2 * beta(elso,"resl") + alpha(elso,"resl") * adjust(elso,"resl")) *
```

ac(elso,"resl"))) +
(ac("slssl","openl") * ((ac("slss1","resl") + ac("slss1","openl")/2) *
beta("slssl","openl") + alpha("slssl","openl")*adjust("slssl","openl"))) +
(ac("slss2","agrl") * ((ac("slss2","resl") + ac("slss2","agrl")/2) *
beta("slss2","agrl") + alpha("s1ss2","agrl")*adjust("s1ss2","agrl"))) +
(ac("s2ssl","openl") * ((ac("s2ssl","resl") + ac("s2ssl","openl")/2) *
beta("s2ss1","open1") + alpha("s2ss1","open1") * adjust("s2ss1","open1"))) +

```
(ac("s2ss2","agr1") * ((ac("s2ss2","res1") + ac("s2ss2","agr1")/2) *
beta("s2ss2","agrl") + alpha("s2ss2","agr1") * adjust("s2ss2","agr1"))) +
(ac("s3ss1","agrl") * ((ac("s3ss1","resl") + ac("s3ss1","agrl")/2) *
beta("s3ss1","agrl") + alpha("s3ss1","agr1")* adjust("s3ss1","agrl"))) +
(ac("s3ss2","agr1")* ((ac("s3ss2","res1") + ac("s3ss2","agr1")/2) *
beta("s3ss2","agr1") + alpha("s3ss2","agr1") * adjust("s3ss2","agr1"))) +
(ac("slss1","agr1")* ((ac("s1ss1","res1") + ac("s1ss1","open1") +
ac("s1ss1","agr1")/2) * beta("s1ss1","agr1") + alpha("s1ss1","agrl") *
adjust("s1ss1","agrl"))) +
(ac("s1ss2","open1") * ((ac("s1ss2","res1") + ac("s1ss2","agr1") +
ac("s1ss2","open1")/2) * beta("s1ss2","open1") + alpha("s1 ss2","open1") *
adjust("s1ss2","open1"))) +
```

(ac("s2ss1","agrl") * ((ac("s2ss1","res1") + ac("s2ss1","open1") +
ac("s2ss1","agrl")/2) * beta("s2ss1","agr1") + alpha("s2ss1","agr1") *
adjust("s2ss1","agrl"))) +
(ac("s2ss2","open1") * ((ac("s2ss2","res1") + ac("s2ss2","agr1") +
ac("s2ss2","open1")/2) * beta("s2ss2","open1") + alpha("s2ss2","open1") *
adjust("s2ss2","open1"))) +
(ac("s3ss1","open1") * ((ac("s3ss1","res1") + ac("s3ss1","agr1") +
ac("s3ss1","open1")/2) * beta("s3ss1","open1") + alpha("s3ss1","open1") *
adjust("s3ss1","openl"))) +
(ac("s3ss2","open1") * ((ac("s3ss2","res1") + ac("s3ss2","agr1") +
ac("s3ss2","open1")/2) * beta("s3ss2","open1") + alpha("s3ss2","open1") *
adjust("s3ssl","openl"))) +
sum(elso,(ac(elso,"f40") * (((ac(elso,"res1") + ac(elso,"openl") + ac(elso,"agrl") +
(ac(elso,"f40")+
ac(elso,"f45") $+\mathrm{ac}($ elso,"f50" $)+\mathrm{ac}($ elso,"f60" $)+\mathrm{ac}($ elso,"f80" $)+\mathrm{ac}($ elso,"f120") +
ac(elso,"park"))/2) * beta(elso,"f40") + alpha(elso,"f40") * adjust(elso,"f40"))))) +
sum(elso,(ac(elso,"f45") * ((ac(elso,"resl") + ac(elso,"openl") + ac(elso,"agrl") + (ac(elso,"f40")+
ac(elso,"f45")+ ac(elso,"f50") + ac(elso,"f60") + ac(elso,"f80") + ac(elso,"f120") + ac(elso,"park"))/2) * beta(elso,"f45") + alpha(elso,"f45") * adjust(elso,"f45"))))) +
sum(elso,(ac(elso,"f50") * (((ac(elso,"res1") + ac(elso,"open1") + ac(elso,"agrl") + (ac(elso,"f40")+
ac(elso,"f45")+ ac(elso,"f50") + ac(elso,"f60") $+\mathrm{ac}($ elso,"f80") $+\mathrm{ac}(\mathrm{elso}, " \mathrm{f} 120$ ") + ac(elso,"park"))/2) * beta(elso,"f50") + alpha(elso,"f50") * adjust(elso,"f50"))))) +
sum(elso,(ac(elso,"f60") * (((ac(elso,"res1") + ac(elso,"open1") + ac(elso,"agr1") + (ac(elso,"f40")+
ac(elso,"f45")+ ac(elso,"f50") + ac(elso,"f60") + ac(elso,"f80") + ac(elso,"f120") +

```
ac(elso,"park"))/2) * beta(elso,"f60") + alpha(elso,"f60") * adjust(elso,"f60"))))) +
sum(elso,(ac(elso,"f80") * (((ac(elso,"res1") + ac(elso,"open1") + ac(elso,"agr1") +
(ac(elso,"f40")+
ac(elso,"f45")+ ac(elso,"f50") + ac(elso,"f60") + ac(elso,"f80") + ac(elso,"f120") +
ac(elso,"park"))/2) * beta(elso,"f80") + alpha(elso,"f80") * adjust(elso,"f80"))))) +
sum(elso,(ac(elso,"f120") * (((ac(elso,"res1") + ac(elso,"openl") + ac(elso,"agr1") +
(ac(elso,"f40")+
ac(elso,"f45")+ ac(elso,"f50") + ac(elso,"f60") + ac(elso,"f80") + ac(elso,"f120") +
ac(elso,"park"))/2) * beta(elso,"f120") + alpha(elso,"f120") * adjust(elso,"f120"))))) +
sum(elso,(ac(elso,"park") * (((ac(elso,"res1") + ac(elso,"open1") + ac(elso,"agrl") +
(ac(elso,"f40")+
ac(elso,"f45")+ ac(elso,"f50") + ac(elso,"f60") + ac(elso,"f80") + ac(elso,"f120") +
ac(elso,"park"))/2) * beta(elso,"park") + alpha(elso,"park") * adjust(elso,"park")))));
```

$\mathrm{NLV}=(\operatorname{sum}($ elso, (adac2.1(elso,"res1")/2 * beta(elso,"res1") + alpha(elso,"res1") * adjust(elso,"res1"))
*
adac2.l(elso,"res1"))) +
(adac2.1("s1ss1","open1") * ((adac2.1("s1ss1","res1") + adac2.1("s1ss1","open1")/2) *
beta("s1ss1","open1") + alpha("s1ss1","open1")*adjust("s1ss1","open1"))) +
(adac2.l("s1ss2","agr1") * ((adac2.l("s1ss2","res1") + adac2.l("s1ss2","agr1")/2) *
beta("s1ss2","agr1") + alpha("s1ss2","agr1")*adjust("s1ss2","agr1"))) +
(adac2.1("s2ss1","open1") * ((adac2.1("s2ss1","res1") + adac2.1("s2ss1","open1")/2) *
beta("s2ss1","open1") + alpha("s2ss1","open1") * adjust("s2ss1","open1"))) +
(adac2.l("s2ss2","agr1") * ((adac2.l("s2ss2","res1") + adac2.l("s2ss2","agr1")/2) *
beta("s2ss2","agr1") + alpha("s2ss2","agr1") * adjust("s2ss2","agr1"))) +
(adac2.l("s3ss1","agr1") * ((adac2.1("s3ss1","res1") + adac2.l("s3ss1","agr1")/2) *
beta("s3ss1","agr1") + alpha("s3ss1","agr1") * adjust("s3ss1","agr1"))) +
(adac2.1("s3ss2","agr1") * ((adac2.1("s3ss2","res1") + adac2.l("s3ss2","agr1")/2) *
beta("s3ss2","agr1") + alpha("s3ss2","agr1") * adjust("s3ss2","agr1"))) +
(adac2.1("s1ss1","agr1") * ((adac2.l("s1ss1","res1") + adac2.1("s1ss1","open1") +
adac2.1("s1ss1","agr1")/2) * beta("s1ss1","agr1") + alpha("s1ss1","agr1") *
adjust("s1ss1","agr1"))) +
(adac2.1("s1ss2","open1") * ((adac2.1("s1ss2","res1") + adac2.1("s1ss2","agr1") +
adac2.l("s1ss2","open1")/2) * beta("s1ss2","open1") + alpha("s1ss2","open1") *
adjust("s1ss2","open1"))) +
(adac2.1("s2ss1","agr1") * ((adac2.l("s2ss1","res1") + adac2.1("s2ss1","open1") +
adac2.1("s2ss1","agr1")/2) * beta("s2ss1","agr1") + alpha("s2ss1","agr1") *
adjust("s2ss1","agr1"))) +
(adac2.1("s2ss2","open1") * ((adac2.1("s2ss2","res1") + adac2.1("s2ss2","agr1") +
adac2.1("s2ss2","open1")/2) * beta("s2ss2","open1") + alpha("s2ss2","open1") *
adjust("s2ss2","open1"))) +

```
(adac2.l("s3ssl","openl") * ((adac2.l("s3ss1","resl") + adac2.l("s3ssl","agrl") +
adac2.l("s3ssl","openl")/2) * beta("s3ssl","openl") + alpha("s3ssl","openl") *
adjust("s3ssl","openl"))) +
(adac2.1("s3ss2","openl") * ((adac2.1("s3ss2","res1") + adac2.1("s3ss2","agr1") +
adac2.l("s3ss2","open1")/2) * beta("s3ss2","open1") + alpha("s3ss2","openl") *
adjust("s3ss1","openl"))) +
```

sum(elso,(adac2.1(elso,"f40") * (((adac2.1(elso,"res1") + adac2.1(elso,"open1") + adac2.1(elso,"agrl") + (adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.l(elso,"f80") + adac2.1(elso,"f120") +
adac2.l(elso,"park"))/2) * beta(elso,"f40") + alpha(elso,"f40") * adjust(elso,"f40"))))) +

```
sum(elso,(adac2.1(elso,"f45") * (((adac2.1(elso,"res1") + adac2.1(elso,"open1") + adac2.1(elso,"agrl") +
(adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.1(elso,"f80") +
adac2.l(elso,"f120") +
adac2.l(elso,"park"))/2) * beta(elso,"f45") + alpha(elso,"f45") * adjust(elso,"f45"))))) +
sum(elso,(adac2.1(elso,"f50") * (((adac2.1(elso,"res1") + adac2.1(elso,"openl") + adac2.1(elso,"agrl") +
(adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.1(elso,"f80") +
adac2.1(elso,"f120") +
adac2.1(elso,"park"))/2) * beta(elso,"f50") + alpha(elso,"f50") * adjust(elso,"f50"))))) +
```

sum(elso,(adac2.1(elso,"f60") * (((adac2.1(elso,"res1") + adac2.1(elso,"open1") + adac2.1(elso,"agr1") +
(adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.1(elso,"f80") +
adac2.1(elso,"f120") +
adac2.l(elso,"park"))/2) * beta(elso,"f60") + alpha(elso,"f60") * adjust(elso,"f60"))))) +
sum(elso,(adac2.1(elso,"f80") * (((adac2.1(elso,"res1") + adac2.1(elso,"open1") + adac2.1(elso,"agrl") +
(adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.1(elso,"f80") +
adac2.1(elso,"f120") +
adac2.1(elso,"park"))/2) * beta(elso,"f80") + alpha(elso,"f80") * adjust(elso,"f80"))))) +
sum(elso,(adac2.1(elso,"f120") * (((adac2.1(elso,"res1") + adac2.1(elso,"open1") + adac2.1(elso,"agr1")
$+$
(adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.1(elso,"f80") +
adac2.1(elso,"f120") +
adac2.1(elso,"park"))/2) * beta(elso,"f120") + alpha(elso,"f120") * adjust(elso,"f120"))))) +
sum(elso,(adac2.1(elso,"park") * (((adac2.1(elso,"res1") + adac2.1(elso,"open1") + adac2.1(elso,"agr1")
$+$
(adac2.1(elso,"f40")+
adac2.1(elso,"f45")+ adac2.1(elso,"f50") + adac2.1(elso,"f60") + adac2.1(elso,"f80") +
adac2.1(elso,"f120") +
adac2.1(elso,"park"))/2) * beta(elso,"park") + alpha(elso,"park") * adjust(elso,"park")))));
$\mathrm{OBIO}=\mathrm{SUM}(\mathrm{S}, \mathrm{WT}(\mathrm{S}) /(1+\mathrm{EXP}(-\mathrm{ALPH}-\mathrm{BET} *$


AC1(ELSO,HAB)))))));

```
EXPD = SUM(S,WT(S)/(1+EXP(-ALPH-BET *
LOG(.01 + (1/THETA) *
(
(spac(s,"res")* sum(elso,adac2.1(elso,"res1")))+
(spac(s,"open")* sum(elso, adac2.1(elso,"open1"))) +
(spac(s,"agr") * sum(elso, adac2.l(elso,"agr1"))) +
(spac(s,"ncon") * sum(elso,((adac2.1(elso,"f40") + .888*adac2.l(elso,"f45") + .8*adac2.1(elso,"f50")+
.6666*adac2.1(elso,"f60") + .5 * adac2.1(elso,"f80") + . 3333 * adac2.1(elso,"f120")))) +
(spac(s,"mcon") * sum(elso,((.5*adac2.1(elso,"f80") + .111* adac2.1(elso,"f45") +
.2 * adac2.1(elso,"f50") + .333* adac2.1(elso,"f60") + .333* adac2.1(elso,"f120")))) +
(spac(s,"ocon") * sum(elso, ((.333 * adac2.1(elso,"f120") + adac2.l(elso,"park"))))
))\)\)))
;
NBIO = EXPD * BIOPRICE;
AB(S) = SUM(HAB,SPAC(S,HAB) * SUM(ELSO,AC1(ELSO,HAB)));
NAB(S)=(spac(s,"res")* sum(elso,adac2.1(elso,"res1")))+
(spac(s,"open") * sum(elso, adac2.1(elso,"open1"))) +
(spac(s,"agr") * sum(elso, adac2.1(elso,"agr1"))) +
(spac(s,"ncon") * sum(elso,((adac2.1(elso,"f40") + .888*adac2.1(elso,"f45") + .8*adac2.1(elso,"f50")+
.6666*adac2.1(elso,"f60") + .5 * adac2.1(elso,"f80") + .3333 * adac2.1(elso,"f120"))))) +
(spac(s,"mcon") * sum(elso,((.5*adac2.l(elso,"f80") + .111* adac2.l(elso,"f45") +
.2* adac2.1(elso,"f50") + .333* adac2.1(elso,"f60") + .333 * adac2.l(elso,"f120")))) +
(spac(s,"ocon") * sum(elso, ((.333 * adac2.1(elso,"f120") + adac2.l(elso,"park"))));
```

FILE RES / k: \shunkn|gamsloutput 11000 .out /
PUT RES;
PUT /"MPRICE = " MPRICE//;
PUT /"BIODIVERSITY PRICE =" BIOPRICE//;
PUT /"ORIGINAL MARKET VALUE =" OLV//;
PUT /"NEW MARKET VALUE =" NLV//;
PUT /"ORIGINAL BIODIVERSITY VALUE $=$ " OBIO//;
PUT /"NEW BIODIVERSITY VALUE =" NBIO//;
PUT /"EXPECTED BIODIVERSITY INDEX =" EXPD//;

## PUT

"SPECIES ORIG ABUNDANCE NEW ABUNDANCE"//;
LOOP (S, PUT S.TL:18, AB(S):18:1 NAB(S):18:3/;
);

PUT /"ADJUSTED AREAS"//;
PUT" f40 f45 f50"//;
LOOP (ELSO, PUT ELSO.TL:9:1, ADAC2.L(ELSO,'f40'):15:1, ADAC2.L(ELSO,'f45'):15:1,ADAC2.L(ELSO,'f50'):15:1/;
);
PUT /"ADJUSTED AREAS CONTINUED"//;
PUT " f60 f80 fl20"//;
LOOP (ELSO, PUT ELSO.TL:9:1,ADAC2.L(ELSO,'f60'):15:1, ADAC2.L(ELSO,'f80'):15:1,ADAC2.L(ELSO,'f120'):15:1/;
);

PUT /"ADJUSTED AREAS CONTINUED"//;
PUT " park open agr"//;
LOOP (ELSO, PUT ELSO.TL:9:1,ADAC2.L(ELSO,'park'):15:1, ADAC2.L(ELSO,'openl'):15:1,ADAC2.L(ELSO,'agr1'):15:1/;
);
PUT /"ADJUSTED AREAS CONTINUED"//;
PUT " res "//;
LOOP (ELSO, PUT ELSO.TL:9:1,ADAC2.L(ELSO,'res1'):15:1/; ;

## Appendix 7.

Total opportunity cost curve for biodiversity
Marginal cost curve, imperiled species biodiversity.



## Appendix 8.

Population index values, $\mathrm{P}_{\mathrm{s}}$, for seven land allocation opportunities.

|  |  | Original Allocation Allocation | 2025 <br> Allocation | $\mathrm{B}=20 \mathrm{mil}$ Allocation | Max Bio Allocation | Imperiled $B=20 \mathrm{mil}$ Allocation | Imperiled Max Bio Allocation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Ambystoma | ile | 4998.5 | 2199.296 | 5338.008 | 5655.947 | 3371.515 | 2839.609 |
| Ambystoma | macrodactyl | 5001.8 | 4960.951 | 5302.039 | 5288.609 | 4952.11 | 5029.55 |
| Aneides | ferreus | 4999.5 | 2438.724 | 5660.755 | 6153.024 | 4752.401 | 4002.641 |
| Ensatina | eschscholtzii | 5001.2 | 3738.416 | 4647.034 | 4861.387 | 2708.077 | 2326.877 |
| Plethodon | dunni | 5001.7 | 1819.121 | 4755.076 | 5115.057 | 3576.257 | 3012.05 |
| Plethodon | vehiculum | 5000.4 | 2271.647 | 4490.024 | 4841.404 | 3491.072 | 2940.305 |
| Taricha | granulosa | 4999.9 | 4051.518 | 5043.68 | 5228.723 | 3493.493 | 3265.797 |
| Dicamptodon | tenebrosus | 1499.6 | 581.098 | 1467.933 | 1568.302 | 1023.709 | 862.204 |
| Ascaphus | truei | 1500.4 | 168.928 | 1773.038 | 1925.202 | 1443.654 | 1215.896 |
| Pseudacris | regilla | 5000.9 | 4637.928 | 5406.409 | 5389.26 | 4468.216 | 4415.999 |
| Rana | aurora | 1501.2 | 892.5 | 1588.243 | 1652.506 | 1227.793 | 1166.305 |
| Ardea | herodias | 5001.1 | 3051.368 | 6414.303 | 6547.193 | 4854.42 | 4713 |
| Butorides | virescens | 4999.6 | 3822.128 | 4753.41 | 4948.11 | 2427.013 | 2044.116 |
| Branta | canadensis | 151.3 | 159.794 | 172.3 | 166.602 | 268.436 | 281.611 |
| Aix | sponsa | 5000.4 | 4463.485 | 7857.8 | 7660.78 | 0 | 0 |
| Anas | Platyrhynch | 4999.4 | 5040.59 | 5694.303 | 5726.625 | 9032.594 | 9887.091 |
| Lophodytes | Cucullatus | 4999.8 | 3136.818 | 7709.753 | 7807.914 | 2140.075 | 1802.447 |
| Cathartes | Aura | 5001.4 | 5974.824 | 5030.705 | 5127.933 | 3969.405 | 3979.456 |
| Pandion | Haliaetus | 5000 | 2345.113 | 13819.53 | 14173.22 | 13902.26 | 10286.03 |
| Elanus | Leucurus | 625 | 621.963 | 742.584 | 743.37 | 1001.023 | 1095.721 |
| Haliaeetus | Leucocephal | 1499.8 | 143.443 | 2672.221 | 2927.627 | 2367.234 | 1993.768 |
| Circus | Cyaneus | 5000.2 | 6570.204 | 4821.802 | 4812.602 | 6614.235 | 7239.951 |
| Accipiter | striatus | 4998.9 | 3265.116 | 4412.832 | 4585.466 | 3049.745 | 2737.073 |
| Accipiter | cooperii | 5000.8 | 3159.728 | 4966.665 | 5087.128 | 3016.239 | 2599.318 |
| Accipiter | gentilis | 624.6 | 0 | 690.468 | 765.156 | 675.499 | 568.929 |
| Buteo | jamaicensis | 5000.6 | 5433.006 | 5257.06 | 5280.15 | 4167.36 | 4171.318 |
| Aquila | chrysaetos | 5001.6 | 8041.699 | 4833.824 | 4959.832 | 4242.115 | 4204.42 |
| Falco | sparverius | 5000.9 | 6298.065 | 5340.6 | 5083.004 | 6068.407 | 6215.913 |
| Dendragapus | obscurus | 4999.8 | 3774.993 | 4362.014 | 4652.22 | 3083.083 | 2596.682 |
| Bonasa | umbellus | 4999.2 | 4538.515 | 4414.421 | 4537.45 | 1866.588 | 1572.106 |
| Oreortyx | pictus | 1499 | 2606.987 | 1187.592 | 1153.323 | 397.603 | 435.216 |
| Charadrius | vociferus | 5000.3 | 6042.832 | 5635.347 | 4756.385 | 8549.089 | 7680.288 |
| Gallinago | gallinago | 4999.4 | 5040.59 | 5694.303 | 5726.625 | 9032.594 | 9887.091 |
| Brachyramph | marmoratus | 149.9 | 0 | 496.465 | 550.168 | 485.701 | 409.075 |
| Columba | fasciata | 4999.2 | 3698.318 | 4354.327 | 4542.033 | 2307.65 | 1944.765 |
| Zenaida | macroura | 4999.9 | 7233.75 | 4614.522 | 4468.743 | 5081.657 | 5365.961 |
| Tyto | alba | 5001.1 | 5291.052 | 6101.815 | 5785.299 | 8555.647 | 8532.473 |
| Otus | kennicottii | 4999.7 | 3884.378 | 5121.88 | 5253.149 | 2617.098 | 2107.05 |
| Bubo | virginianus | 4998.8 | 3889.508 | 5545.961 | 5678.844 | 4555.331 | 4364.57 |
| Glaucidium | gnoma | 1500.4 | 754.429 | 1475.501 | 1577.535 | 1046.126 | 881.084 |
| Athene | cunicularia | 150.2 | 151.4 | 171.035 | 172.006 | 271.305 | 296.971 |
| Strix | occidentalis | 624.6 | 0 | 919.606 | 1019.08 | 899.668 | 757.733 |
| Strix | varia | 5000.1 | 1652.873 | 7275.659 | 7674.952 | 4271.183 | 3597.342 |
| Asio | otus | 1500.3 | 1519.186 | 1187.623 | 1192.84 | 306.366 | 258.032 |
| Asio | flammeus | 1499 | 1511.394 | 1707.406 | 1717.098 | 2708.375 | 2964.591 |


|  |  | Original Allocation Allocation | $2025$ <br> Allocation | 20 million Allocation | Max Bio Allocation | Imperiled 20 million Allocation | Imperiled Max Bio Allocation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| A | aca | 1499.5 | 647.927 | 158 | 1685.478 | 1029.687 | 867.239 |
| Chordeiles | minor | 5000.7 | 9795.762 | 2848.807 | 2769.533 | 2221.896 | 2432.091 |
| Chaetura | va | 5000 | 2345.113 | 13819.53 | 14173.22 | 13902.26 | 10286.03 |
| Calypte |  | 1500.4 | 1515.402 | 2273.145 | 2080.024 | 238.786 | 0 |
| Selasphorus | rufus | 5000.5 | 5497.079 | 4103.183 | 4087.816 | 1850.571 | 93.372 |
| Melanerpes | lewis | 1500.1 | 1339.046 | 2357.34 | 2298.234 | 0 | 0 |
| Melanerpes | formicivorus | 626.6 | 559.636 | 983.572 | 959.001 | 4.678 | 12 |
| Sphyrapicus | ruber | 1500.8 | 871.699 | 1575.584 | 1606.396 | 862.802 | 73.4 |
| Picoides | pubescens | 1500.6 | 1935.009 | 1879.343 | 1756.33 | 124.458 | 0 |
| Picoides | villosus | 1499.7 | 689.045 | 1680.897 | 1783.53 | 1080.499 | 10.034 |
| Colaptes | auratus | 5000.5 | 4744.237 | 5115.674 | 4948.721 | 3871.414 | 3337.148 |
| Dryocopus | pileatus | 1500.1 | 625.085 | 1616.627 | 1734.702 | 1183.617 | 996.884 |
| Contopus | borealis | 1500.1 | 902.347 | 1409.922 | 1527.02 | 1156.717 | 974.228 |
| Contopus | sordidulus | 1500.4 | 1649.327 | 1641.949 | 1576.061 | 428.514 | 268.101 |
| Empidonax | traillii | 1500.2 | 3390.989 | 587.266 | 552.411 | 0 | 0 |
| Empidonax | hammondii | 1499.5 | 1270.443 | 837.625 | 888.345 | 581.347 | 489.631 |
| Empidonax | difficilis | 1500.8 | 644.584 | 1525.545 | 1607.509 | 966.701 | 803.045 |
| Tyrannus | verticalis | 4998.9 | 4989.297 | 5883.914 | 5896.067 | 8237.389 | 9016.658 |
| Eremophila | alpestris | 4999.4 | 5040.59 | 5694.303 | 5726.625 | 9032.594 | 9887.091 |
| Progn | subis | 625.2 | 564.452 | 1343.26 | 1472.71 | 1222.473 | 1029.61 |
| Tachycineta | bicolor | 5000.9 | 5657.461 | 5299.831 | 5367.176 | 2492.694 | 2353.24 |
| Tachycineta | thalassina | 5001.5 | 9700.62 | 3183.458 | 2682.877 | 2700.469 | 2160.721 |
| Hirundo | pyrrhonota | 5000.1 | 9524.51 | 5426.52 | 1378.756 | 6861.744 | 0 |
| Hirundo | rustica | 5000.6 | 6431.078 | 5612.451 | 4380.432 | 8361.678 | 6825.216 |
| Cyanocitta | stelleri | 4999.6 | 4464.121 | 4203.031 | 4448.569 | 2754.301 | 2319.769 |
| Aphelocoma | californica | 4999.2 | 5395.92 | 7128.554 | 6320.462 | 2487.297 | 1464.375 |
| Corvus | brachyrhync | 5000.9 | 5868.487 | 5628.817 | 5244.679 | 7145.272 | 7122.187 |
| Corvus | corax | 1500.6 | 733.575 | 1689.352 | 1810.361 | 1491.238 | 1336.246 |
| Parus | atricapillus | 5000 | 5660.663 | 6140.553 | 5690.851 | 1205.836 | 482.079 |
| Parus | rufescens | 000.8 | 3246.337 | 4151.979 | 4412.667 | 3262.111 | 2673.462 |
| Psaltriparus | minimus | 4999.7 | 8612.215 | 4276.699 | 3932.381 | 935.032 | 619.543 |
| Sitta | canadensis | 5000.1 | 2407.33 | 4560.697 | 4842.622 | 3477.125 | 2849.679 |
| Sitta | carolinensis | 1499.7 | 1381.954 | 2335.835 | 2243.797 | 58.715 | 0 |
| Certhia | americana | 1499.6 | 544.77 | 1755.949 | 1855.32 | 1142.333 | 949.054 |
| Thryomanes | bewickii | 1501.5 | 2330.628 | 1254.557 | 1193.614 | 135.238 | 107.524 |
| Troglodytes | aedon | 1498.9 | 2334.392 | 1318.359 | 1222.781 | 349.891 | 211.91 |
| Troglodytes | troglodytes | 1499.3 | 1244.311 | 1320.836 | 1395.842 | 841.384 | 708.644 |
| Regulus | satrapa | 1500.4 | 754.429 | 1475.501 | 1577.535 | 1046.126 | 881.084 |
| Sialia | mexicana | 1498.5 | 2269.275 | 1351.694 | 1303.37 | 1210.715 | 1269.808 |
| Catharus | ustulatus | 5000.6 | 4360.209 | 4009.628 | 4200.628 | 2305.961 | 1942.162 |
| Turdus | migratorius | 5001.1 | 5497.596 | 4660.798 | 4584.637 | 3751.153 | 3600.248 |
| Ixoreus | naevius | 1499.8 | 177.666 | 1750.231 | 1897.878 | 1406.292 | 1184.429 |
| Chamaea | fasciata | 4999.7 | 10201.66 | 2905.496 | 2776.343 | 0 | 0 |
| Bombycilla | cedrorum | 5001.3 | 7577.527 | 5138.829 | 4645.308 | 1794.966 | 1269.808 |
| Vireo | solitarius | 1498.9 | 1270.751 | 1558.658 | 1563.925 | 354.188 | 298.31 |

Original 202520 million Max Bio Imperiled Imperiled Allocation Allocation Allocation Allocation 20 million Max Bio Allocation

Allocation Allocation

| Vireo | huttoni |
| ---: | ---: |
| Vireo | gilvus |
| Vermivora | celata |
| Dendroica | petechia |
| Dendroica | coronata |
| Dendroica | nigrescens |
| Dendroica | occidentalis |
| Oporornis | tolmiei |
| Geothlypis | trichas |
| Wilsonia | pusilla |
| Iteria | virens |
| Piranga | ludoviciana |
| Pheucticus | melanoceph |
| Passerina | amoena |
| Pipilo | maculatus |
| Spizella | passerina |
| Poecetes | gramineus |
| Passerculus | sandwichen |
| Ammodramus | savannarum |
| Melospiza | melodia |
| Zonotrichia | leucophrys |
| Junco | hyemalis |
| Agelaius | phoeniceus |
| Sturnella | neglecta |
| Euphagus | cyanocephal |
| Molothrus | ater |
| Icterus | bullockii |
| Carpodacus | purpureus |
| Carpodacus | mexicanus |
| Loxia | curvirostra |
| Carduelis | pinus |
| Carduelis | psaltria |
| Carduelis | tristis |
| Coccothrauste | vespertinus |
| Sorex | vagrans |
| Sorex | pacificus |
| Sorex | bendirii |
| Sorex | trowbridgii |
| Sorex | sonomae |
| Neurotrichus | gibbsii |
| Scapanus | townsendii |
| Scapanus | orarius |
| Myotis | lucifugus |
| Myotis | yumanensis |
| Myotis | evotis |
|  |  |


| 1499.6 | 616.381 | 1467.137 | 1526.736 | 717.344 | 604.172 |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 4999.6 | 6180.349 | 6476.007 | 6094.662 | 359.11 | 0 |
| 5000.1 | 8189.659 | 4660.53 | 4470.579 | 64.917 | 0 |
| 1500.1 | 1339.046 | 2357.34 | 2298.234 | 0 | 0 |
| 5000.4 | 544.647 | 5585.886 | 6062.352 | 4526.737 | 3812.578 |
| 5000.4 | 4243.184 | 5222.778 | 5238.715 | 1173.156 | 988.073 |
| 1500.4 | 536.136 | 1416.289 | 1526.167 | 1084.982 | 913.81 |
| 1500.6 | 3008.022 | 918.433 | 879.035 | 0 | 0 |
| 5000.2 | 4888.042 | 6268.776 | 6240.268 | 6637.623 | 7265.552 |
| 4999.2 | 5897.299 | 2080.772 | 1997.551 | 0 | 0 |
| 1500.1 | 1339.46 | 2357.34 | 2298.234 | 0 | 0 |
| 5000.3 | 2409.694 | 4747.295 | 4996.331 | 2766.256 | 2329.839 |
| 4999.8 | 2686.719 | 5773.9 | 5950.744 | 2400.112 | 2021.459 |
| 1498.8 | 1383.4 | 2185.151 | 2144.145 | 711.007 | 778.269 |
| 1501 | 2198.818 | 1342.416 | 1257.054 | 458.022 | 404.495 |
| 1500 | 1975.551 | 1683.105 | 1597.608 | 918.372 | 901.154 |
| 1499.8 | 1498.354 | 1760.191 | 1764.388 | 2493.202 | 2729.062 |
| 5000.9 | 5029.68 | 5742.604 | 5769.997 | 8840.809 | 9677.163 |
| 149.6 | 147.869 | 181.284 | 181.091 | 224.528 | 245.769 |
| 5001.3 | 6911.8 | 4267.832 | 4067.503 | 3008.177 | 2900.883 |
| 4999.6 | 7977.346 | 4346.537 | 4101.737 | 3323.803 | 3323.005 |
| 5001.6 | 4626.5 | 3605.728 | 3631.591 | 2343.066 | 2174.443 |
| 4999.9 | 5019.815 | 5774.632 | 5798.502 | 8700.479 | 9523.557 |
| 1501.4 | 1503.039 | 1750.327 | 1755.769 | 2544.656 | 2785.384 |
| 4999.4 | 5471.419 | 5799.762 | 5393.46 | 8249.615 | 8248.629 |
| 5000.4 | 5731.503 | 4729.265 | 4637.901 | 3821.601 | 3831.089 |
| 1499.5 | 2209.212 | 1656.488 | 1503.897 | 182.139 | 0 |
| 1500.7 | 1148.414 | 1259.216 | 1310.253 | 645.609 | 543.755 |
| 5001.4 | 5554.37 | 6565.964 | 5812.992 | 4848.232 | 4039.831 |
| 1499.5 | 390.022 | 1472.692 | 1588.264 | 1129.816 | 951.571 |
| 4999.6 | 3592.023 | 4320.761 | 4639.758 | 3284.836 | 2766.605 |
| 1500.7 | 1484.852 | 2022.995 | 1947.554 | 1358.953 | 1392.692 |
| 1500.7 | 2129.118 | 1498.555 | 1429.295 | 1176.032 | 1198.125 |
| 4999.6 | 1592.13 | 5687.539 | 6138.52 | 4402.696 | 3708.107 |
| 1500.3 | 1471.447 | 1566.363 | 1574.749 | 1343.411 | 1361.049 |
| 623.3 | 594.846 | 574.864 | 591.325 | 246.587 | 207.684 |
| 1499.1 | 1439.072 | 1293.899 | 1342.479 | 647.104 | 545.014 |
| 1500.7 | 1443.966 | 1259.575 | 1311.823 | 669.521 | 563.894 |
| 5000.2 | 4781.458 | 4510.043 | 4652.381 | 2039.946 | 1718.115 |
| 1500.3 | 1144.016 | 1634.643 | 1704.011 | 844.373 | 711.161 |
| 1499.7 | 1548.983 | 1726.922 | 1669.264 | 2010.642 | 2058.209 |
| 4999.3 | 4823.307 | 4537.437 | 4678.13 | 2961.359 | 2771.575 |
| 1498.7 | 1307.127 | 1605.872 | 1613.155 | 1495.253 | 1385.067 |
| 625.8 | 501.366 | 799.674 | 820.134 | 725.128 | 659.987 |
| 627 | 614.211 | 571.723 | 583.115 | 427.774 | 401.334 |
|  |  |  |  |  |  |


|  |  | Original Allocation Allocation | $2025$ <br> Allocation | 20 million Allocation | Max Bio Allocation | Imperiled 20 million Allocation | Imperiled Max Bio Allocation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Myotis | thysanodes | 627.3 | 469.231 | 668.352 | 703.042 | 562.801 | 524 |
| Myotis | volans | 623.4 | 312.776 | 900.717 | 961.836 | 811.854 | 740.435 |
| Myotis | californicus | 1499.5 | 1163.397 | 1635.195 | 1698.8 | 1375.636 | 1293.285 |
| ionycteris | noctivagans | 1499.4 | 1403.591 | 1900.817 | 1970.052 | 1639.978 | 14 |
| Eptesicus | fuscus | 1500.6 | 1463.469 | 1869.516 | 1889.113 | 1802.441 | 1662. |
| Lasiurus | cinereus | 1499.7 | 917.844 | 2179.667 | 2275.813 | 1752.634 | 1562.025 |
| Plecotus | townsendii | 1500.8 | 1280.033 | 1436.193 | 1455.221 | 1134.58 | 1030.857 |
| Antrozous | pallidus | 624.8 | 664.035 | 781.434 | 728.925 | 797.761 | 783.389 |
| Sylvilagus | bachmani | 5000.4 | 4803.681 | 4804.025 | 4917.943 | 2895.781 | 2744.676 |
| Lepus | americanus | 1499.9 | 1653.545 | 1097.428 | 1125.049 | 467.768 | 393.9 |
| Lepus | californicus | 1499.9 | 1507.066 | 1727.777 | 1735.417 | 2628.854 | 2877.548 |
| Aplodontia | rufa | 1500.6 | 1562.216 | 947.193 | 968.044 | 561.501 | 524.857 |
| Tamias | townsendii | 1500.3 | 1493.754 | 1312.874 | 1366.031 | 687.454 | 578.998 |
| Spermophilus | beecheyi | 5000.3 | 5157.387 | 5943.522 | 5779.812 | 7861.279 | 8294.71 |
| Sciurus | griseus | 1498.5 | 538.582 | 1760.908 | 1811.163 | 899.012 | 723.74 |
| Tamiasciurus | douglasii | 5000.2 | 609.451 | 5473.889 | 5923.039 | 4305.556 | 3626 |
| Glaucomys | sabrinus | 1499.9 | 359.326 | 1606.717 | 1670.262 | 1147.247 | 903.74 |
| Thomomys | mazama | 4998.2 | 6162.439 | 2502.264 | 2442.685 | 1660.575 | 1817.668 |
| Thomomys | bulbivorus | 1499.9 | 1559.57 | 1740.569 | 1694.733 | 2533.463 | 2677.86 |
| Castor | canadensis | 5000.6 | 4930.471 | 4854.695 | 4906.94 | 4240.121 | 4178.39 |
| Peromyscus | maniculatus | 5000.1 | 4888.549 | 4754.134 | 4829.912 | 3574.998 | 3429.4 |
| Neotoma | fuscipes | 1500.1 | 1339.046 | 2357.34 | 2298.234 | 0 |  |
| Neotoma | cinerea | 4999.7 | 4796.851 | 4342.48 | 4501.743 | 2141.57 | 1803.706 |
| Clethrionomys | californicus | 1500 | 820.371 | 1593.502 | 1693.97 | 1056.587 | 889.895 |
| Phenacomys | albipes | 626.3 | 676.315 | 477.787 | 498.75 | 269.004 | 226.56 |
| Phenacomys | longicaudus | 1500.2 | 0 | 2208.888 | 2447.822 | 2160.998 | 1820.06 |
| Microtus | townsendii | 1500.8 | 1477.029 | 1844.492 | 1839.865 | 2147.054 | 2350.168 |
| Microtus | longicaudus | 5000.3 | 10083.35 | 3008.922 | 2878.342 | 0 |  |
| Microtus | oregoni | 1499.7 | 1805.668 | 944.684 | 959.329 | 352.694 | 297.051 |
| Microtus | canicaudus | 4998.9 | 4973.701 | 5942.079 | 5948.058 | 7994.15 | 8750.408 |
| Zapus | trinotatus | 1500.1 | 978.816 | 1576.727 | 1662.661 | 951.975 | 801.78 |
| Erethizon | dorsatum | 5000.8 | 4810.079 | 4432.915 | 4583.708 | 2515.051 | 2244.57 |
| Canis | latrans | 4999.3 | 5204.854 | 4703.496 | 4704.198 | 4155.956 | 4127.605 |
| Vulpes | vulpes | 1500 | 1613.286 | 1572.054 | 1547.256 | 1515.496 | 1593.616 |
| Urocyon | cinereoarge | 1499.9 | 1326.75 | 1381.757 | 1443.447 | 754.705 | 635.639 |
| Ursus | americanus | 1499.2 | 1446.788 | 1357.852 | 1394.091 | 752.013 | 664.976 |
| Procyon | lotor | 4999.9 | 5001.246 | 4654.517 | 4624.69 | 3126.964 | 2695.393 |
| Martes | americana | 624.9 | 541.624 | 578.37 | 610.537 | 363.155 | 305.86 |
| Martes | pennanti | 149.8 | 105.836 | 139.592 | 149.765 | 104.613 | 88.108 |
| Mustela | erminea | 4999.7 | 4811.727 | 4604.877 | 4738.629 | 2804.783 | 2605.468 |
| Mustela | frenata | 4999.7 | 4811.727 | 4604.877 | 4738.629 | 2804.783 | 2605.468 |
| Spilogale | gracilis | 1499.3 | 1623.093 | 1300.006 | 1332.804 | 538.008 | 453.129 |
| Mephitis | mephitis | 4999.6 | 6597.104 | 5722.794 | 5574.678 | 4607.876 | 4499.84 |
| Felis | concolor | 1500.4 | 1629.618 | 1425.639 | 1440.5 | 1183.657 | 1190.482 |
| Lynx | rufus | 1500.5 | 1479.325 | 1363.537 | 1383.722 | 918.165 | 871.66 |


|  |  | Original Allocation Allocation | $2025$ <br> Allocation | 20 million Allocation | Max Bio Allocation | Imperiled 20 million Allocation | Imperiled Max Bio Allocation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Cervus | elaphus | 5000.3 | 5369.89 | 4975.793 | 5049.404 | 4919.169 | 5074.584 |
| Odocoileus | hemionus | 4997.9 | 5440.558 | 4813.339 | 4822.916 | 4280.424 | 4287.761 |
| Chrysemys | picta | 150.3 | 149.999 | 176.801 | 177.174 | 247.917 | 271.37 |
| Clemmys | marmorata | 150.4 | 150.425 | 175.904 | 176.39 | 252.595 | 276.49 |
| Elgaria | coerulea | 5000 | 6435.796 | 3897.16 | 3908.301 | 2199.728 | 1912.243 |
| Elgaria | multicarinata | 4999.2 | 6830.35 | 5448.118 | 5155.213 | 2803.753 | 2713.702 |
| Sceloporus | occidentalis | 4999.8 | 8138.1 | 2785.028 | 2545.251 | 239.381 | 15.361 |
| Eumeces | skiltonianus | 5000.4 | 8332.108 | 4689.414 | 4180.147 | 676.924 | 51.202 |
| Charina | bottae | 1498.5 | 1923.064 | 1326.215 | 1301.599 | 736.791 | 663.7 |
| Coluber | constrictor | 1499.3 | 1510.235 | 1595.241 | 1564.789 | 1379.843 | 1343.598 |
| Contia | tenuis | 624.1 | 662.125 | 651.795 | 641.54 | 361.084 | 332.106 |
| Diadophis | punctatus | 1501.1 | 1493.474 | 1537.589 | 1518.151 | 1042.024 | 958.131 |
| Pituophis | catenifer | 4999.7 | 5926.449 | 5293.099 | 5061.158 | 5481.481 | 5470.877 |
| Thamnophis | ordinoides | 4999.9 | 5714.739 | 4313.718 | 4197.46 | 3496.069 | 3363.272 |
| Thamnophis | sirtalis | 4999.7 | 5451.373 | 4962.224 | 4838.208 | 4630.627 | 4454.147 |
| Crotalus | viridis | 1501.2 | 1421.407 | 2054.474 | 2027.626 | 1272.328 | 1392.692 |

