AN ASSESSMENT OF STREAM FISH VULNERABILITY AND AN

EVALUATION OF CONSERVATION NETWORKS IN MISSOURI

A Thesis

Presented to

the Faculty of the Graduate School

at the University of Missouri

In Partial Fulfillment

Of the Requirements for the Degree

Master of Science

by

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DECEMBER 2014

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ACKNOWLEDGMENTS

I would first like to thank the United States Geological Service National Climate Change and Wildlife Science Center for funding this project. I would also like to thank the Missouri Department of Conservation (MDC) for providing the fish community data which served as the foundation upon which this project was completed. Specifically, I would like to thank Matt Combes and Dr. Doug Novinger, who not only provided me with tremendous sources of data for Missouri's stream fish communities, but also shared with me their expertise and knowledge by reviewing my work and offering invaluable insights. Dorothy Butler of MDC also generously provided fish records from the Missouri Natural Heritage Database. I would also like to thank Gust Annis and the Missouri Resource Assessment Partnership for providing me with GIS data without which this project would not have been possible. Dennis Figg of MDC also provided me with both insight and data related to Missouri's conservation networks. Bob Hrabik and Amy Davis, both of MDC, provided valuable feedback and insight throughout the development of this thesis. I would also like to thank those who attended the Missouri Conservation Network Evaluation Meeting we organized at the MDC Central Regional Office in April of 2014. We were honored to have 29 participants from a variety of groups including MDC, US Fish and Wildlife Service, US National Parks Service, and the Nature Conservancy attend and provide valuable feedback. I would also like to thank Dr. Yin-Phan Tsang and Dr. Dana Infante from Michigan State University for providing information on species responses to climate change as well as feedback on my work.

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There have been many people throughout this process who have contributed greatly to both my personal and professional growth, while giving me great memories and lots of laughs. I'd like to thank my fellow graduate students and post docs in the Department of Fisheries and Wildlife Sciences, specifically Landon Pierce, Jason Harris, Jeff Fore, Pablo Oliero, Dr. Jacob Westhoff, Emily Pherigo, and Andy Dinges, for their mentorship, thoughtfulness, friendship, and for keeping me sane during long hours in the basement. I would like to thank my parents Randy and Sandy Sievert for always encouraging me to learn and to follow my passions, and my fiancé Veronica Conaway for her support, encouragement, and patience throughout this process.

I would also like to thank Dr. Tim Matisziw and Dr. Michelle Staudinger for providing me with unique perspectives and helpful feedback throughout the development of my thesis. Dr. Jodi Whittier was instrumental to this project. She was always available to share her knowledge and experience with me, without which I would have a lot less hair on top of my head and this project would not have been as successful. Most of all I would like to thank my advisor, Dr. Craig Paukert, for doing everything he could to give me an opportunity to be successful. Craig's guidance helped me grow as a thinker, professional, researcher, writer, and person. I could not have asked for a more supportive, accessible, and good-natured advisor. I owe so much gratitude to so many people; this thesis is truly a product of the efforts and encouragement of all who have supported me.

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AN ASSESSMENT OF STREAM FISH VULNERABILITY AND AN EVALUATION OF CONSERVATION NETWORKS IN MISSOURI

Nicholas A. Sievert

Dr. Craig P. Paukert, Thesis Supervisor

ABSTRACT

The conservation of aquatic biodiversity is largely dependent on the ability of researchers and managers to identify vulnerable species and prioritize conservation actions. Stream fish are facing tremendous threats due to habitat degradation, stream network fragmentation, and climate change. For this project two vulnerability indices were developed to quantify the relative vulnerability of stream fish in Missouri. These indices allowed us to identify the most vulnerable stream fish in Missouri, and also to compare how the use of traits differed from the use of measured species responses in quantifying vulnerability. The most valuable areas for stream fish conservation within and complementary to Missouri's conservation networks were identified through the use of a systematic conservation planning approach. Species representation, weighting based on vulnerability and listing status, and species-specific responses to upstream habitat degradation were utilized to identify priority areas with the conservation planning tool Zonation. This information can assist managers in identifying the species most in need of conservation and where the best opportunities exist for taking management actions. The frameworks for assessing both vulnerability and conservation value can be used or adapted for addressing similar concerns in other regions.

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DESCRIPTION OF CHAPTERS

The two chapters of this thesis were written as independent manuscripts which will be submitted to peer-reviewed journals. For this thesis we included a general introduction and conclusion to tie the elements of the two chapters together into a more complete narrative. Because each chapter was written as an independent manuscript there is some overlap in the information conveyed in the introductions, as well as in the study area and data sections of the materials and methods, and references, tables, and figures that were included separately for each chapter. A single set of appendices were generated for the entire thesis. Intended co-authors of the manuscripts were listed after each chapter title and the thesis was written using plural nouns to include co-authors.

GENERAL INTRODUCTION

Lotic ecosystems are among the most imperiled on the planet (Allan and Flecker 1993, Abell 2002, Dudgeon et al. 2006). Freshwater biodiversity is declining at a faster rate than in any terrestrial biome (Ricciardi and Rasmussen 1999), and the threats posed by anthropogenic land-use, climate change, flow modification, and invasive species are likely to exacerbate the situation (Ricciardi and Rasmussen 1999, Dudgeon et al. 2006, Staudt et al. 2013). A combined extinction rate for unionid mussels, crayfishes, fish, and amphibians is near 4% per decade, which is significantly higher than that of other taxa (Ricciardi and Rasmussen 1999). Throughout most of the world, information on the status of biodiversity is limited, biased, or absent (Dudgeon et al. 2006). However, in areas with robust data, researchers are finding alarming trends in the number of imperiled species. In North America approximately 39% of aquatic species are considered imperiled (Jelks et al. 2008) and in regions with less data, such as Asia, the pressures of dense human populations, rapid growth, and high levels of biodiversity and endemics likely present an even bleaker scenario (Dudgeon *et al.* 2006). This evidence suggests the conservation of aquatic biodiversity likely requires conservation and management actions, or substantial biodiversity losses are likely. In Missouri, the status of aquatic biodiversity is following similar trends. Of the 210 native fish species, 66 (31%) have been designated as species of conservation concern (Sowa et al. 2007, Missouri Natural Heritage Program 2012). The long-term conservation of aquatic

biodiversity depends on our ability to identify threats, determine species vulnerability, and develop effective conservation strategies.

Lotic biodiversity is expected to be influenced by a number of threats including warming stream temperatures, alterations to flow regimes, and habitat degradation across the globe (Allan and Flecker 1993, Eaton and Scheller 1996, Poff et al. 2002, Allan 2004). Stream temperatures are expected to increase by approximately 3.6 degrees Celsius by 2095 based on an intermediate (A1B) estimate of average annual air temperature and air to stream temperature conversions (Eaton and Scheller 1996, Girvetz et al. 2009), which has been predicted to result in species extirpations and range contractions (Eaton and Scheller 1996, Mohseni et al. 2003, Lyons et al. 2010). Even if temperature shifts do not result in range loss, they have the potential to affect the performance of populations by influencing energetics, growth, and feeding (Pease and Paukert 2013, Westhoff and Paukert 2014). Alterations to stream flow regimes from changes in the timing, amount, and type (snow vs rain) of precipitation are expected to impact stream fish (Poff et al. 2002). Some stream fish species depend upon predictable seasonal flow patterns for successful reproduction, and changes to these patterns could result in a decline of reproductive success (Poff et al. 2002). Increases in the intensity of rainfall on fewer rain days are predicted, leading to an increased magnitude and frequency of flooding, which has been linked to shifts in species composition and local species extirpations (Poff et al. 1997, 2002). Summer and fall stream flow may be reduced in many regions where increased frequency and magnitudes of droughts have been predicted (Poff et al. 2002). Species which require consistent or high amounts of

discharge may be lost from stream segments impacted by drought conditions (Larimore *et al.* 1959, Stanley *et al.* 1997). Habitat alteration and degradation resulting from anthropogenic land use impacts is another major threat to stream fish (Malmqvist and Rundle 2002). Continued agricultural and urban development and the modification of streams and rivers through channelization, dredging, and damming, are expected to continue to have negative impacts on sensitive stream fish species (Malmqvist and Rundle 2002, Allan 2004).

The state of Missouri is heavily impacted by both urban and agricultural development. Approximately 37% of Missouri is cropland (which includes both harvested and pastured lands), and while urban land comprises only 3.7% of Missouri's area, it is highly concentrated in the St. Louis and Kansas City areas (Blodgett and Lea 2005). Land use varies substantially by region, with agriculture and grassland dominating in the north, forests in the south and urban around St. Louis and Kansas City (Blodgett and Lea 2005). Identifying areas for protection of aquatic biodiversity is important as urban and agricultural land use encompass about 40% of the state (Blodgett and Lea 2005).

In order to assess the status of aquatic biodiversity conservation in Missouri, we addressed two primary questions:

- How vulnerable are Missouri species and communities to current and future threats?
- 2: How well is aquatic biodiversity represented within Missouri's conservation networks and which areas are the most valuable for stream fish conservation?

We can prioritize conservation efforts to benefit the species and communities in greatest need by developing an understanding of which species are vulnerable to climate change and habitat degradation and identifying areas which are important to the conservation of those species. We evaluated aquatic species representation within Missouri's established conservation networks using species distribution models and identified priority areas for stream fish conservation using systematic conservation planning techniques. These networks include the existing conservation network (ECN) composed of areas established for the primary purpose of wildlife conservation, Conservation Opportunity Areas (COAs) which are areas established in the Missouri Comprehensive Wildlife Strategy (Hoskins 2005) as target areas for conservation action, and Priority Watersheds (PWs) which were established as a part of the Missouri Department of Conservation Fisheries Program as target areas for conservation and management actions (Corson et al. 2010). This allowed for the evaluation of how well aquatic biodiversity is represented within these networks, and a ranking of stream segments within and complementary to each established network based on their value for stream fish conservation.

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CHAPTER 1

VULNERABILITY ASSESSMENT FOR MISSOURI STREAM FISH TO CLIMATE CHANGE AND HABITAT DEGRADATION

Nicholas Sievert and Dr. Craig P. Paukert

ABSTRACT

Understanding the future impacts and threats of climate and land use change are critical for long-term biodiversity conservation. We developed and compared two indices to assess the vulnerability of stream fish in Missouri. Both indices assessed vulnerability based on species environmental tolerances, rarity, range size, dispersal ability, and the average connectivity of the streams occupied by each species. These indices differed in how the environmental tolerance components were classified, specifically vulnerability to habitat degradation, warming stream temperatures, and alterations to flow regimes. In one index environmental tolerance components were classified based on measured species responses, while in the second index environmental tolerance was classified based on species traits. We also determined 1) if vulnerability scores were consistent between indices, 2) the relationship between species conservation status (species listed as Missouri species of conservation concern) and vulnerability score, and 3) how species vulnerable to these three threats were spatially distributed. Vulnerability scores were calculated for all 133 species with the trait association index, while only 99 species were able to be evaluated using the species response index, because a number of species lacked response data. The range and

mean for scores from the trait association index were greater than those from the species response index, largely due to the species response index's inability to evaluate many rare species, which generally have high vulnerability scores for the trait association index. The indices were consistent in classifying vulnerability to habitat degradation, but showed substantial variation for vulnerability to increases in stream temperature and alterations to flow regimes. This is likely because the current climate change impacts have had a minimal measurable influence on stream fish communities, limiting the opportunities to observe species-specific responses to these threats, while traits can be more generally linked to vulnerability. Both indices showed higher mean vulnerability scores for listed species than unlisted species, which provided a coarse measure of validation. The distribution of vulnerable species in Missouri showed consistent patterns between indices, with the Ozark subregion generally having higher numbers and proportions of vulnerable species per site than in other subregions, which suggests that if conservation actions target habitats rather than species, both indices will perform similarly. This vulnerability assessment provides valuable information regarding the status of Missouri stream fish species and we believe this framework can be used, and updated, for the development of vulnerability assessments in other regions.

INTRODUCTION

Stream ecosystems have some of the most imperiled communities on Earth and freshwater biodiversity is declining at a higher rate than most other taxa groups (Allan and Flecker 1993; Abell 2002; Dudgeon et al. 2006). Continuing habitat degradation, warming stream temperatures, and alterations to flow regimes will likely cause continuing declines (Ricciardi and Rasmussen 1999; Dudgeon et al. 2006). Conservation actions aimed at protecting aquatic biodiversity are critical for preventing future biodiversity losses (Master et al. 1998). In order to plan for long-term biodiversity conservation we must gain a better understanding of how impacts and threats, such as climate change and habitat degradation, affect aquatic species. Conservation and management of stream fish species will require researchers and managers to identify which species are vulnerable, or likely to experience harm, under future conditions (Turner et al. 2003; Glick et al. 2011; Poff et al. 2012). This study developed a new vulnerability assessment framework that identifies species vulnerable to warming stream temperatures, alterations in flow regimes, and habitat degradation. These are among the primary threats identified as drivers of future stream fish declines (Malmqvist and Rundle 2002; Poff et al. 2002).

Climate change, which is expected to increase stream temperatures and alter flow regimes, is one of the most significant threats facing stream fish (Eaton and Scheller 1996; Poff *et al.* 2002). In Missouri, the average annual air temperature is expected to rise approximately 4 degrees Celsius by 2095 based on an intermediate (A1B) climate scenario (Girvetz *et al.* 2009). This change in air temperature is predicted

to increase Missouri stream temperatures by approximately 3.6 degrees Celsius (Eaton and Scheller 1996), although this relationship is complicated locally by factors such as groundwater input, riparian shading, and stream size (Mohseni and Stefan 1999). In Wisconsin, a 3 degrees Celsius air temperature increase is predicted to result in the loss of 343,034 km of stream habitat for cool- and cold-water fishes, which includes a species extirpation (Lyons et al. 2010). Eaton and Scheller (1996) found that the amount of habitat available for cold- and cool-water fish species would be reduced nationwide by approximately 50% with a doubling of atmospheric CO₂, resulting in species extirpations in many streams and dramatic range contractions for many species. In another study of streams across the United States cold-water fish habitat was predicted to decrease by 36% and cool-water fish habitat by 15% with a doubling of atmospheric CO₂ concentrations (Mohseni et al. 2003). Warm-water streams in the Great Plains sometimes achieve maximum temperatures which are at or near the physiological limits of some fish species, and an increase in stream temperature of just a few degrees is predicted to result in local extirpation and even extinction of some species (Matthews and Zimmerman 1990). Even in cases where stream fish are not extirpated from a stream segment, research has shown that some species are likely to be subject to decreases in fitness based on changes in energetics and growth (Pease and Paukert 2013; Westhoff and Paukert 2014). These studies provide strong evidence which suggests some species of stream fish will decline or face local extirpations as stream temperatures warm.

Changes to flow regimes, related to the seasonal timing and amount of precipitation and snowmelt, are expected to have substantial impacts on stream fish (Poff et al. 2002). Increases in the intensity of rainfall on fewer rain days are predicted, which may lead to an increased magnitude and frequency of flooding; this has been linked to shifts in species composition and local species extirpations (Poff et al. 1997, 2002). Summer and fall stream flow may be reduced in many regions because of increased evapotranspiration and decreased precipitation frequency, earlier snowmelt could reduce spring flows, and increased frequency and magnitudes of droughts and floods are expected to occur (Poff et al. 2002; Wuebbles and Hayhoe 2004). Stream fish vary in their ability to cope with prolonged periods of low flow, and species which require high levels of discharge are likely to be eliminated from many sites (Larimore et al. 1959; Stanley et al. 1997). The variability of precipitation patterns is likely to increase, which may cause declines in species which exhibit equilibrium or periodic life history strategies (Poff et al. 2002; Olden and Kennard 2010; Mims and Olden 2012, 2013).

Another major threat to lotic biodiversity is habitat alteration and degradation (Malmqvist and Rundle 2002). Anthropogenic modifications including the conversion of land to agricultural or urban uses or the modification of streams and rivers from channelization, dredging, and damming often result in altered and degraded stream conditions (Malmqvist and Rundle 2002; Allan 2004). Degraded streams have been associated with losses of aquatic biodiversity (Allan 2004). Indices of biological integrity (IBIs) have identified many stream fish species as intolerant of habitat degradation

(Esselman *et al.* 2011). Additionally, life history traits such as lithophilic spawning, and benthic invertivory have been used as criteria for IBIs measuring habitat degradation, because species with those traits are sensitive to degradation (Berkman and Rabeni 1987; Barbour *et al.* 1999; Simon 1999).

With this knowledge of threats facing stream fish species the vulnerability of species can be assessed. For the purposes of this thesis, vulnerability is defined as the extent to which a species is likely to be negatively impacted by the cumulative effects of climate change and habitat degradation (Turner et al. 2003; Schnieder et al. 2007; Glick et al. 2011). In order to conduct a vulnerability assessment for stream fish species, a framework which accounts for the impacts of multiple threats is needed. Vulnerability is often determined using a framework that assesses a species sensitivity, exposure, and adaptive capacity to threats (Turner et al. 2003; IPCC 2007; Glick et al. 2011a; Poff et al. 2012; Staudinger et al. 2013). A number of organizations, government agencies, and researchers have developed vulnerability assessments using this framework (Bagne et al. 2011; Glick et al. 2011; Young et al. 2011). Two prominent vulnerability assessment tools, the System for Assessing Vulnerability of Species (SAVS) developed by the United States Forest Service, and the NatureServe Climate Change Vulnerability Index (NSCCVI) developed for NatureServe, were applied to Missouri stream fish (Bagne et al. 2011; Young *et al.* 2011). The applicability of these vulnerability assessments to stream fish species in Missouri to be limited. In Missouri accurate models of current and future stream temperature and flow have not yet been developed. The absence of these models limits our ability to assess the exposure component of these assessments. Both

assessment techniques use air temperature models to predict changes in thermal habitat, however in stream systems the relationship between air and water temperatures is confounded by riparian shading, groundwater contribution, and stream size (Mohseni and Stefan 1999; Allan 2004; Caissie 2006; Whitledge et al. 2006; Webb et al. 2008; Westhoff and Paukert 2014). Because this study is only able to express exposure in the most general terms (warming is expected to occur, flow regimes are expected to become less predictable and more extreme events are expected to occur) it could not be used to differentiate species and therefore cannot be included as a scoring component, at least not until adequate models have been developed. Additionally, these tools were designed for use with a wide array of taxa over large spatial scales, which limits their ability to account for some of the nuances of stream fish vulnerability. Another issue with the application of these tools to stream fish is that they depend on information which is largely unknown for stream fish species; examples include knowledge of a species reliance on interspecific interactions, measures of genetic variation, occurrence of bottlenecks in recent evolutionary history, and phenological response to changing seasonal temperature and precipitation dynamics (Bagne et al. 2011; Young *et al.* 2011). Currently, there simply is not enough information available to broadly apply these assessment tools to stream fish in a meaningful way. The inability to reasonably apply available vulnerability assessment tools to determine the vulnerability of Missouri's stream fish necessitates the development of a new methodology.

Poff et al. (2012) developed a framework for assessing the threat posed by climate change to freshwater diversity. This framework is a function of three components: exposure to the threats, sensitivity of species to impacts of threats, and freshwater resilience or ability to adapt to or cope with the threats (Table 1). Although this framework deviates from the definitions and structure presented by Glick et al. (2011) which is widely used for developing vulnerability assessments, we believe that it provides a better mechanism for assessing stream fish vulnerability based on the information that is currently available, and therefore this thesis conformed to structure and definitions found in Poff et al. (2012). Species environmental tolerances, which are often expressed as sensitivity in the vulnerability assessment literature, specifically to habitat degradation, stream temperature warming, and alterations to flow regimes, as well as factors such as a species range, rarity, dispersal ability, and the hydrological connectivity of a species habitat can be incorporated into this framework to create a method for assessing the vulnerability of stream fish species. This analysis of species vulnerability will focus on the sensitivity and freshwater resilience components of the Poff et al. (2012) framework as adequate information is not yet available to assess exposure in Missouri.

Species tolerances of habitat degradation, altered flow regimes, and increasing stream temperatures have been assessed using two different approaches; species trait associations (Angermeier 1995; Parent and Schriml 1995; Poff 1997; Olden *et al.* 2007, 2008; Culp *et al.* 2011; Mims and Olden 2013) and measured species responses (Hering *et al.* 2006; Lyons *et al.* 2010, Tsang and Infante, Michigan State University, Unpublished

data). For this thesis two separate indices were developed, one which scores environmental tolerance based on traits and the other which scores based on species responses. The same scoring framework was used for both indices. The trait association approach to classifying environmental tolerance is based on traits which have been linked to vulnerability to habitat degradation, altered flow regimes, and increases in stream temperature in peer-reviewed literature. The species response approach is based on species-specific observations of sensitivity to each of the environmental tolerance components. Species environmental tolerances provide the foundation for the sensitivity component of this vulnerability assessment framework.

Another aspect of the sensitivity component of the vulnerability assessment framework is species dispersal ability (Poff *et al.* 2012). The ability of a fish species to disperse throughout a stream network is critical to the persistence of species faced with threats (Albanese *et al.* 2009; Poff *et al.* 2012). Dispersal, a one-way movement from a site (Lidicker Jr. and Stenseth 1992), may benefit species facing threats by allowing 1) for gene flow between subpopulations (Hanski 1998; Heggenes *et al.* 2006), 2) increased colonization of newly available habitats (Detenbeck *et al.* 1992; Albanese *et al.* 2009), and 3) species to shift their ranges as environmental changes occur at the landscape scale (Matthews and Zimmerman 1990; Tonn 1990).

Both range size and population size can influence the vulnerability of a species (Poff *et al.* 2012). Species with smaller ranges are limited in their ability to withstand stochastic environmental and demographic fluctuations and are therefore more

vulnerable (Angermeier 1995). Species constrained to a small range are more likely to lack access to refugia or an adequate gradient of conditions necessary to cope with the impacts of threats. Rare species, even those which are widely distributed across the landscape, may have increased vulnerability to both deterministic threats such as habitat loss, as well as stochastic events such as invasions and epidemics (Mace *et al.* 2008). Species with narrowly restricted distributions and rare species generally have high levels of vulnerability.

Freshwater resilience, defined as the hydrological connectivity of the landscape, is another component of the vulnerability assessment framework developed by Poff *et al.* (2012). A stream system with high levels of connectivity provides species the opportunity to emigrate to suitable habitats as conditions change (Poff *et al.* 2012). Not only is dispersal difficult because of the restrictive, branching nature of stream networks, but humans have increased the level of fragmentation through the creation of barriers such as dams and road crossings (Jackson and Marmulla 2001; Fagan *et al.* 2002). Species with highly fragmented populations may have an elevated risk of extinction (Fagan *et al.* 2002).

Scientists and decision makers benefit from the ability to identify vulnerable species to make effective decisions regarding the conservation of stream fish. The objectives of this chapter are to 1) classify species' vulnerability based on environmental tolerance to habitat degradation, stream temperature warming, and alterations to flow regimes, dispersal ability, restricted distribution, rarity, and freshwater resilience 2)

compare the use of traits and species responses to identify species environmental tolerance 3) develop indices to measure stream fish vulnerability using our species classifications, 4) determine whether our measures of vulnerability correlate with listing status, and 5) analyze distribution patterns of vulnerable species in Missouri. Our aim is to develop a framework for assessing stream fish vulnerability and to make recommendations for assessing stream fish vulnerability based on an application of our methodology to Missouri's stream fish species.

MATERIALS AND METHODS

Study area

The state of Missouri can be classified into three ecologically unique subregions: the Central Plains, Ozarks, and Mississippi Alluvial Basin (MAB) (Pflieger 1970; Pflieger and Missouri Department of Conservation 1989; Sowa *et al.* 2007) (Figure 1). These subregions feature major differences in geology, landform, soils, land cover, and groundwater influence which create unique habitats and fish communities in the streams of each area (Table 2). The Central Plains, in northern Missouri, is dominated by open grassland and agriculture (Figure 1) with streams associated with wide, gently sloping valleys which have a relatively low gradient, high turbidity, and fine silt and sand substrates (Sowa *et al.* 2007). The Ozarks, in southern Missouri (except the southeastern corner), is a mix of forested areas and open pastureland and agriculture (Sowa *et al.* 2007) (Figure 1). Most of the region has rugged terrain with high relief, which creates higher average stream gradients. Many streams in the Ozarks have substantial groundwater inputs. The substrate of these streams is highly variable but often consists of gravel, cobble, or bedrock with sand and silt in slower moving areas and pools. The MAB is in southeastern Missouri, and is a broad flat plain with low stream gradients, most of which have been channelized for agriculture, which is the predominant land cover in the region (Figure 1). The substrate varies throughout the subregion but generally the larger and slower moving streams are characterized by fine silt while streams with faster flow are comprised of sand and small gravel substrates (Sowa *et al.* 2007).

Fish data

Data detailing fish occurrence records were provided by the Missouri Department of Conservation (MDC) from the Resource Assessment and Monitoring Program (RAM) database. This study used 1499 fish samples which were collected using a statewide stratified random sampling design (Fischer and Combes 2003) of unique stream segments collected between April and September, 2000 to 2011 from permanent wadeable streams (2nd-5th Strahler order) (Figure 2). All samples within the RAM database were collected using standardized methods with block nets enclosing a sample area which was then thoroughly sampled via electrofishing and seining (Fischer and Combes 2003), with all individual specimens identified to species and enumerated.

Scoring system and classification of species vulnerability to threats

Species vulnerability scores were calculated using two different approaches: one based on species responses and the other linked to species traits (Figure 3). Species

vulnerability was classified and scored for both indices according to species environmental tolerance, dispersal ability, restricted distribution, rarity, and freshwater resilience. Species environmental tolerance scores were calculated by summing the scores for habitat, temperatures, and flow vulnerability separately for each of the two indices, while all other criteria were scored the same way for both indices (Figure 3). Species sensitivity scores were calculated by multiplying rarity scores by the sum of environmental tolerance, dispersal ability, and range size scores, while the final species score is the sum of the sensitivity and freshwater resilience scores (Figure 3). Scores were assigned to each species based on the classification of species vulnerability to each threat category. Rarity was incorporated as a multiplicative factor rather than an additive component because rarity is expected to compound the impacts of the other components. Scores for each threat category were assigned as either discrete (0 or 1) for vulnerable or not classifications, continuous-additive (0 to 1) for vulnerability which was quantified along a gradient and incorporated into the vulnerability index as an additive component, or continuous-multiplicative (1 to 2) for vulnerability which was quantified along a gradient and incorporated into the vulnerability index as a multiplicative component. Multiplicative components were scaled from 1 to 2, rather than 0 to 1, because these components are only intended to have no effect or increase a species score, not reduce it.

Environmental tolerance was scored using the discrete value system by summing the vulnerable (1) or not (0) classifications for vulnerability to habitat degradation, warming stream temperatures, and alterations to flow regimes (Figure 3). Two separate

environmental tolerance scores were calculated; one using a traits-based approach and another using a measured species response approach. Classifications for the traitsbased approach were made after a thorough review of the literature identified traits linked to vulnerability to each of the threats. Lithophilic spawning and benthic invertivory were used to classify species as vulnerable to habitat degradation as both have been linked to sensitivity to sedimentation, a frequent product of habitat degradation (Berkman and Rabeni 1987; Mims *et al.* 2010). Species which exhibit either of these traits were classified as vulnerable to habitat degradation. Species classified as cool- or cold-water adapted have been identified as vulnerable to warming stream temperatures and were therefore classified as vulnerable to warming stream temperatures in this assessment (Matthews and Zimmerman 1990; Eaton and Scheller 1996; Mohseni et al. 2003; Lyons et al. 2010; Mims et al. 2010). As stream flows become less predictable, and incidences of both extreme droughts and floods increase (Poff et al. 2002), species exhibiting the periodic life history strategy (prefer predictable flow patterns), and the equilibrium life history strategy (prefer low variability in flow patterns) are expected to be more vulnerable than species exhibiting the opportunistic life history strategy (thrive in streams with unpredictable and variable flow patterns) (Winemiller and Rose 1992; Olden and Kennard 2010; Mims and Olden 2012, 2013).

The second index used species responses to identify the environmental tolerance of species to habitat degradation, stream temperature warming, and alterations to flow regimes (Figure 3). A substantial amount of research has been conducted to identify species which are intolerant of habitat degradation, particularly

due to the proliferation of IBIs (Karr 1981; Karr *et al.* 1986; Lyons *et al.* 1996; Karr and Chu 1997; Whittier *et al.* 2007; Pont *et al.* 2009). Esselman *et al.* (2011) conducted a literature review of fish based IBIs from across North America and compiled a list of intolerant species. Species which have been classified as intolerant within the World Wildlife Fund ecoregions (Abell *et al.* 2008) which intersect Missouri (ecoregions numbered 143, 146, 147, 148, 149) were classified as vulnerable to habitat degradation for our species response index. Tsang *et al.* (Michigan State University, Unpublished data) used stream gauge station temperature and flow data to identify species which decreased in abundance with temperature and flow metrics corresponding to future climate change predictions. Species demonstrating decreased abundance in streams with flow metrics associated with climate change predictions were classified as vulnerable to predicted alterations to flow regimes. Species showing decreased abundance in streams with temperature metrics associated with climate change predictions were classified as vulnerable to warming stream temperatures.

Species ability to disperse throughout the stream network is critical to recolonization after disturbance, establishment in new habitat made available via changes to the environment, and maintenance of healthy metapopulation dynamics (Albanese *et al.* 2009; Campbell Grant 2011; Radinger and Wolter 2013). Species with limited dispersal abilities are expected to be vulnerable (Poff *et al.* 2012). Radinger *et al.* (2013) quantified the average dispersal distance for the mobile component (dispersers) for nine fish families. Species from three families (Fundulidae, Cottidae, and Percidae), which had an average dispersal distance of less than 250 meters (Radinger

and Wolter 2013), were classified as vulnerable based on their limited dispersal abilities and received scores of 1 (Figure 3). Species from all other families received scores of 0 for the dispersal ability component of these vulnerability indices.

Species range size and rarity have been linked to vulnerability due to environmental change (Poff et al. 2012). Range size was scored using the continuousadditive approach (Figure 3). In order to quantify vulnerability based on range size the number of ecological drainage unites (EDUs) in which a species has a known occurrence were counted. These values were then scaled from 0 for species which occur in all 17 EDUs within Missouri to 1 for species which occur in a single EDU. Intermediate values were calculated along this scale; for instance if a species was found in 25% of EDUs it would receive a score of 0.75, if a species was found in 88% of drainages it would receive a score of 0.12. Similarly, species rarity was calculated by using scaled values from 1 to 2. First, a cutoff for which species would be considered common was needed. This was achieved by examining a histogram of the percent of sites occupied by each species. A threshold was identified at less than 10% of sites (87 species; Figure 4). Using the continuous-multiplicative approach, species which were found at <10% of sites received scores scaled from 1 (10% of sites) to 2 (A single site, 0.01% of sites). Through this scaling system a species found at seven percent of sites would receive a score of 1.3 while a species found at two percent of sites would receive a score of 1.8.

Freshwater resilience scores were calculated as the ratio of the number of connected stream kilometers per occurrence scaled using the continuous additive

approach from 0 for the maximum value (1,787 km/occurrence) to 1 for the minimum value (123 km/occurrence) (Figure 3). The amount of connected stream kilometers was calculated by partitioning out 2nd-6th order stream segments in the Missouri Resource Assessment Partnership's stream network GIS layer (Blodgett and Lea 2005). Stream segments classified as stream or stream/river (i.e., not reservoir or impoundment) were isolated via a definition query, which clipped each stream network at any location which flowed into an impoundment. The number of connected stream kilometers for each intact stream segment was calculated with the Rivex tool (Hornby 2014). This information allowed us to quantify the number of connected stream kilometers for each species occurrence. This information was used to determine the average number of connected stream kilometers for each species occurrence inhabit areas with relatively high stream network connectivity, while species with relatively low values occur in fragmented habitats with low connectivity.

Index comparisons

Several analyses were conducted to determine whether the species response and trait association approaches for classifying environmental tolerance achieved similar results, whether state-listed species received higher vulnerability scores than unlisted species, and whether there were patterns in the distribution of vulnerable species between aquatic subregions. The percent agreement between the two approaches for classifications of vulnerability to habitat degradation, warming stream temperatures, and alterations to flow regimes was used to determine the consistency of

the approaches to classifying environmental tolerance. A Fischer's exact test was used to determine if the indices showed a positive odd's ratio (α <0.01) for vulnerable classifications of the environmental tolerance components of the two approaches. A two-way analysis of variance was used to determine if mean vulnerability score differed by listing status, Missouri species of conservation concern including all federally threatened and endangered species (Missouri Natural Heritage Program 2012), and environmental tolerance classification approach. Pearson's correlation analysis was used to determine if the two indices were related based on the number and proportion of vulnerable species at each of the study sites for each of the three components of environmental tolerance. An analysis of variance was used to determine whether the mean number and mean proportion of vulnerable species to each of the three environmental tolerance components differed by Missouri's three aquatic subregions. All statistical calculations were performed using R statistical software (R Development Core Team 2011).

RESULTS

Index development

Vulnerability scores for all 133 species were calculated using a trait association approach to classifying environmental tolerance (Appendix 1). The mean vulnerability score using the trait association approach was 3.84 (+/- 0.35 95% Cl). Using the species response approach for classifying environmental tolerance 99 of the 133 species could be evaluated and the mean score for this approach was 2.38 (+/- 0.28 95% Cl) (Appendix

 The remaining 34 species were not evaluated because they lacked the information necessary for classifying environmental tolerances. The trait association approach classified 71, 55, and 54 species vulnerable to habitat degradation, warming stream temperatures, and alterations to flow regimes respectively (Appendix 2). The species response approach classified 43, 3, and 18 species vulnerable to habitat degradation, warming stream temperatures, and alterations to flow regimes respectively (Appendix 3). For each of the 133 species rarity, dispersal, range, and freshwater resilience scores were calculated (Appendix 4). The mean rarity score was 1.48 with 65.4% of species classified as being rare based on occurrence rates of <10% of sites. Thirty four species (25.6%) were classified as vulnerable based on limited dispersal ability. The mean number of EDU's in which a species was present was 8 (of 17 total), which equates to a score of 0.53. The mean value for freshwater resilience was 0.53 which equates to an average of 743 connected kilometers per occurrence.

Index comparisons

The species response and trait association approaches to classifying environmental tolerance showed similarities and differences in classification. The most consistent classification between methods was for vulnerability to habitat degradation (65.7%), followed by warming temperatures (60.1%) and lastly flow regime alteration (46.5%) (Table 3). There was a high degree of positive association between the trait and response based approaches to classifying vulnerability based on habitat degradation (Odd's ratio of 3.68, p=0.002), while no significant association was found between the trait and response based approaches to classifying vulnerability based on

warming temperatures (Odd's ratio of 0.80, p=0.99) or changes in flow regimes (Odd's ratio of 0.49; p= 0.2) (Table 3).

Mean vulnerability scores differed by both listing status (p<0.001), and index (p<0.001), with no interaction between listing status and index (p=0.794). The mean species response score for listed species was 3.35, while unlisted species averaged scores of 2.29; for the trait association approach, listed species averaged scores of 4.84, while unlisted species averaged scores of 3.57 (Figure 5). Scores for rare species (species occurring in less than 10% of sites) were compared to common species and it was found that rare species had significantly higher vulnerability scores for both approaches (Response: Rare=2.8, Not =1.9, P<0.001; Trait: Rare=4.6, Not=2.5, P<0.001).

The number of vulnerable species at each site had the highest correlation between the trait and response-based approaches for habitat degradation (r=0.94), followed by temperature (r=0.37), and flow (r=0.23) (Figure 6). The proportion of vulnerable species at each site had the highest correlation between both indices for habitat degradation (r=0.87), followed by temperature (r=0.15), and finally flow (r=-0.25) (Figure 6). The mean number of vulnerable species per site was greatest in the Ozark subregion for both habitat degradation and temperature using both approaches of classifying environmental tolerances (Table 4). The greatest numbers of species vulnerable per site for flow occurred in the Mississippi Alluvial Basin according to the trait association approach, and the Plains according to the species response approach (Table 4). The mean proportion of vulnerable species per site was highest in the Ozarks

for habitat and temperature for the trait association approach and habitat for the species response approach (Table 4). The greatest proportion of flow vulnerable species per site was highest in the Mississippi Alluvial Basin for the trait association, and in the plains for the species response approach (Table 4).

DISCUSSION

The results of this assessment provide a framework for assessing stream fish vulnerability to climate change and habitat degradation impacts. Stream communities in regions across the globe will be impacted by climate change and habitat degradation (Eaton and Scheller 1996; Poff et al. 2002; Heino et al. 2009), and evaluating species vulnerability will be critical for conservation planning efforts (Poff et al. 2012). This assessment provides a new approach for stream fish vulnerability assessment which can be applied and modified in other regions and improved with further evaluation and refinement. The trait association approach to classifying environmental tolerance was applied to all species while the species response approach could only able be applied to 74 percent of species in this analysis. The trait association approach produced higher vulnerability scores (3.84 versus 2.38), and exhibited a greater range of scores (0-8.75 versus 0-7.82). Both of these discrepancies between the indices largely stems from the inability of the species response index to classify the environmental tolerances of rare species. Of the 34 species for which environmental tolerance could be classified based on measured species responses, 32 were classified as rare (occurring at less than 10% of the study sites). Rare species had significantly higher vulnerability scores than common species using both approaches which likely resulted in higher average and maximum

scores for the trait association index compared to the species response index. Although some regions, such as California, appear to have sufficient information regarding the vulnerability of species to specific metrics, many regions lack the data necessary to elicit the responses of species to environmental threats necessary for a broadly applied vulnerability assessment (Moyle *et al.* 2013). To further assess and quantify the impact of changes in the environment, quantitative assessments of stream fish relationships with potential stressors is needed (Schlosser 1991). Broad scale vulnerability assessments which rely on measured species responses to environmental threats will depend on additional research to determine the impacts of threats on all species. Many of these rare species are likely the most vulnerable, so a failure to include them in a vulnerability assessment would likely leave an important data gap and present a biased assessment.

A traits based approach allows for a more complete assessment of species vulnerability as traits have been described for the vast majority of North American stream fish (Frimpong and Angermeier 2009; Mims *et al.* 2010). The use of traits has flourished for the assessment of stream fish communities, and has proven to be a useful tool in stream fish ecology and conservation research (Parent and Schriml 1995; Poff 1997; Olden and Kennard 2010; Culp *et al.* 2011; McManamay *et al.* 2014). The results presented here suggest a traits based approach allows for a more complete representation of species; while more research linking specific traits to environmental threats would help decrease the uncertainty of these assessments and validate measures of vulnerability.

Substantial discrepancies existed in the number of species classified as vulnerable to environmental tolerance components using the trait and response based approaches. Considering only species evaluated using both the trait association and species response approaches, it was determined that both approaches classified a similar number of species as vulnerable to habitat degradation (47 versus 43 respectively). However, the number of species classified as vulnerable to both warming stream temperatures (38 versus 3) and alterations to flow regimes (47 versus 18) was considerably higher for the trait association approach. This is likely because results derived using traits to classify vulnerability could be extrapolated based on expectations of how species will respond, however when using measured species responses classifications are restricted to observations of responses which have already occurred. Because habitats have already been widely degraded there have been many opportunities to measure species responses in both degraded and intact habitats, however currently observed changes in temperature and flow regimes have only accounted for a small proportion of long-term predictions (Heino et al. 2009), and thus responses to these changes may not yet have taken place. This lack of observations of more extreme conditions predicted in the future has likely led to an inability to classify those species which may show negative responses under future climactic conditions.

Differences were observed in which species classified as vulnerable by the two indices. Vulnerability to habitat degradation was classified relatively consistently by both approaches (65.7% agreement), suggesting that both indices were likely successful in capturing and representing species vulnerability to land use changes. In cases where

the trait association index classified species as vulnerable while the species response index did not, species are often those which have been identified as some of the most tolerant of degradation (e.g. Johnny Darter Etheostoma nigrum, Orangethroat Darter Etheostoma spectabile, and Golden Redhorse Moxostoma erythrurum (Karr 1981)), and other common, tolerant species (e.g. Orangespotted Sunfish Lepomis humilis (Meador and Carlisle 2007), Creek Chub Semotilus atromaculatus (Miller et al. 1988), and Black Bullhead Ameiurus melas (Novomeská and Kováč 2009)). Species classified as vulnerable to habitat degradation by the species response index, which were not by the trait association index, included a number of Centrarchids, (e.g. Warmouth Lepomis qulosus, Longear Sunfish Lepomis megalotis, Smallmouth Bass Micropterus dolomeiu, Shadow Bass Ambloplites ariommus, Ozark Bass Ambloplites constellatus, and Rock Bass Ambloplites rupestris), and some species of the genus Cyprinella (e.g. Whitetail Shiner *Cyprinella galactura*, Spotfin Shiner *Cyprinella spiloptera*, and Steelcolor Shiner *Cyprinella whipplei*). This suggests that although these indices are relatively consistent in classifying vulnerability to habitat degradation, there are still some vulnerable species which were excluded from one approach or the other. The relative consistency between the approaches of classifying habitat vulnerable species suggests that the current understanding of vulnerability to habitat degradation is sufficient for the purposes of a broad-scale assessment using either approach.

In contrast, vulnerability to stream warming and alterations to flow regimes showed major discrepancies between the two approaches. Although only three Missouri stream fish species (White Sucker *Catostomus commersonii*, Smallmouth Bass, and Central Stoneroller *Campostoma anomalum*) have shown measured negative responses to warming stream temperatures, under future conditions many additional species could be vulnerable. There are examples of predicted range contraction for many species under future climate projections, even if that range contraction has yet to be observed (Mohseni *et al.* 2003; Heino *et al.* 2009; Lyons *et al.* 2010). Based on the trait association approach, which allows negative responses to be extrapolated even though they may not yet have been observed, a large number of Cyprinids (14 of 33) and Percids (8 of 15) will likely be vulnerable to future warming.

Discrepancies between the two indices were greatest for flow which suggests that there is a need to resolve the conflicting results between vulnerability predictions based on life history strategy (Mims *et al.* 2010) versus vulnerability predictions based on measured species responses (Tsang and Infante, Michigan State University, Unpublished data). Discrepancies between these indices found a number of Catostomids (12 species), Centrarchids (12 species), and Ictalurids (6 species) had traits linked to vulnerability but showed no negative response, while no species within these families showed a negative response without traits linked to vulnerability. The lack of a measured response for these equilibrium and periodic life history strategists suggests that other factors may complicate the predicted relationship between these traits and decreases in abundance and occupancy associated with changes in stream flow. In contrast, several families had much higher numbers of species showing only negative species responses than only traits linked to vulnerability; including Cyprinids (8 versus 4) and Percids (4 versus 1). Based on the trait research literature it was expected that

these opportunistic species would tolerate or even benefit from projected changes in stream flow regimes (Mims and Olden 2013; McManamay *et al.* 2014), however observations of fish responses to predicted flow metrics show decreases in species abundance (Tsang and Infante, Michigan State University, Unpublished data). These discrepancies suggest that a thoughtful comparison between the response and trait association approaches should be used to select the most appropriate approach, and future research should be conducted to rectify these differences. When selecting an approach for classifying environmental tolerances researchers and managers may need to consider if sufficient information is available for target species, and which approach to classifying environmental tolerance better fits the most current understanding of species vulnerability, particularly regarding the alteration of flow regimes.

Our species vulnerability scores were positively correlated with listing status, which suggests that both of the indices achieve results that align with those of more traditional methods of identifying vulnerable species. In Missouri, species of conservation concern are determined through expert knowledge while also considering rarity, population trends, and threats following the NatureServe Conservation Status Assessment Criteria (Missouri Natural Heritage Program 2012). Using this information as a partial validation it appears that the quantitative assessment of stream fish vulnerability presented here is consistent with expert opinion. All 23 species of conservation concern were able to be evaluated using the trait association index, while only 5 species of conservation concern were with the species response index. Based on the correlations the indices have with species of conservation concern it appears that

species which are currently considered imperiled are likely to be vulnerable to future threats.

Our analysis of the distribution of vulnerable species showed strong spatial trends. Both approaches to classifying environmental tolerance identified the Ozarks as generally having both the highest numbers and proportions of vulnerable species per site than in the other subregions. The exception to this was for flow vulnerability, for which the number and proportion of vulnerable species per site was highest in the MAB according to the trait association approach and the Plains according to the species response approach. Streams in the Ozarks have steeper stream gradients, more diverse substrates, cooler stream temperatures, and less human development than Plains streams and are able to support more sensitive species (Sowa *et al.* 2007). The spatial consistency of the counts and proportions of species classified as vulnerable to habitat degradation and stream warming suggests that management for those threats can be focused on the Ozarks regardless of which approach to classifying environmental tolerance is used. The inconsistency regarding flow vulnerability corroborates the previous comparison of environmental tolerance classification by confirming that further research to determine species vulnerability to predicted alterations to flow regimes is needed. Streams are predicted to experience greater variation in flows (IPCC 2002; Poff et al. 2002), however the impact of increased variation in flow on stream fish is unclear. Decision makers may need to consider which approach to classifying environmental tolerance best fits their understanding of flow vulnerable species when

making decisions regarding the identification and selection of areas for management of these species.

Understanding the impact of uncertainty is crucial for any vulnerability assessment (Patt et al. 2005; Füssel and Klein 2006) and there are many sources of uncertainty that affect our understanding of stream fish vulnerability in Missouri. One of the primary sources of uncertainty in any climate change work is how much the climate will change (Stainforth et al. 2007). Many scenarios have been developed which predict a range of future temperature and precipitation conditions (IPCC 2013). In stream systems this uncertainty is compounded by the fact that there is also uncertainty in how these changes in climate will impact stream conditions (Whitledge et al. 2006; Westhoff and Paukert 2014). In Missouri models do not yet exist which allow us to predict stream temperature and flow conditions now or under predicted future climate scenarios. Although the general trends are known for changes in stream temperature and flow regimes, currently suitable data is not available to differentiate levels of exposure across the Missouri landscape. Because of this exposure, which is often used in vulnerability assessments (Glick et al. 2011; Staudinger et al. 2013), cannot be used as a factor of discrimination in species vulnerability analysis. This assessment is also bound to the uncertainty inherent in predicting the negative effects of environmental threats on species based on both traits and measured responses. Understanding the impact of uncertainty on this assessment is crucial for the interpretation of these results, the applicability of the framework for others, and for providing opportunities to refine the assessment in order to reduce uncertainty when new information becomes available.

There are clear tradeoffs between the two approaches of assessing environmental tolerance. Benefits of a trait association approach include broad applicability to many species, consistency between studies/regions, and ability to link to a causal mechanism (Parent and Schriml 1995; Poff 1997; Frimpong and Angermeier 2009; Culp *et al.* 2011). Benefits of a measured species response approach include direct evidence of species intolerance of threats, and the ability to account for complex interactions between species characteristics and their responses (Williams et al. 2008; Lyons et al. 2010, Tsang and Infante, Michigan State University, Unpublished Data). Disadvantages of a trait association approach include difficulty representing variability in environmental tolerances, limited inference to known and quantified traits, and difficulty accounting for complex species/environmental relationships. Disadvantages of a measured species response approach include restriction to species which have data available (analysis of many rare species are limited by lack of data), limitation of inference to threat impacts which have already occurred in the environment or dependence on laboratory studies to simulate threat impacts, and difficulty isolating the impact of a threat from other factors which may affect a species response. A trait association approach may currently be better suited to assessing stream fish vulnerability because this approach allows us to analyze more species of interest using extrapolations rather than direct observations. Research which investigates species responses to threats could add to the feasibility of a response based assessment of vulnerability. A more complete set of information regarding species thermal and flow tolerances, bioenergetics, and population viability coupled with landscape models of

stream temperatures and flow regimes could lend insight for the development of response based models which could predict species specific vulnerability.

Many stream fish species are already threatened with local extirpation, and increases in habitat degradation, stream warming, and alterations to flow regimes of streams will likely result in increases in the number of species impacted as well as the magnitude of this imperilment (Poff et al. 2012). To more effectively conserve stream fish, decision makers need a better understanding of which species are most vulnerable to each of the threats. This study created a quantifiable index of stream fish vulnerability based on the outline by Poff et al. (2012) that fills this need for the state of Missouri. As more information is gathered for stream fishes the application of more traditional vulnerability assessments following the criteria and framework established by Glick *et al.* (2011) may be possible, however based on currently available information we believe this approach provides a suitable alternative. The assessment of vulnerability presented here allows for identification of species vulnerable to specific threats and also gives us a cumulative measure of a species vulnerability to a suite of threats. This study developed a framework which can serve as a foundation for future vulnerability assessments and can be adapted to include new information regarding species environmental tolerances, additional threats, and the weighting or scale of threats. Our hope is that others can apply and modify this framework for assessment of vulnerable stream fish species in other regions.

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TABLES

Table 1: Vulnerability assessment components used to assess Missouri stream fish vulnerability based on Poff *et al.* 2012.

| Component | Definitions from Poff et al. 2012 |
|--------------------------|--|
| Exposure | Deviation in physico-chemical conditions relative to regional baselines. |
| Sensitivity | Intrinsic factors related to a species environmental tolerance, dispersal ability, genetic adaptation, range, and population size. |
| Freshwater Resilience | Connectivity of aquatic habitats which provides opportunities for species dispersal. |

| | Missouri Land Use (%) | | | | | | |
|-----------|-----------------------|--------|--------|----------------------------|--|--|--|
| | Statewide | Plains | Ozarks | Mississippi Alluvial Basin | | | |
| Urban | 3.7 | 4.1 | 3.5 | 1.8 | | | |
| Cropland | 23.9 | 37.1 | 7.0 | 83.0 | | | |
| Grassland | 33.3 | 35.9 | 34.4 | 3.8 | | | |
| Forest | 34.5 | 17.3 | 51.4 | 5.0 | | | |
| Wetland | 2.1 | 3.5 | 1.0 | 2.5 | | | |
| Water | 2.2 | 2.0 | 2.1 | 3.9 | | | |
| Other | 0.3 | 0.1 | 0.6 | 0.1 | | | |

Table 2: Land use type percentages for the three Missouri aquatic subregions and statewide (Blodgett and Lea 2005).

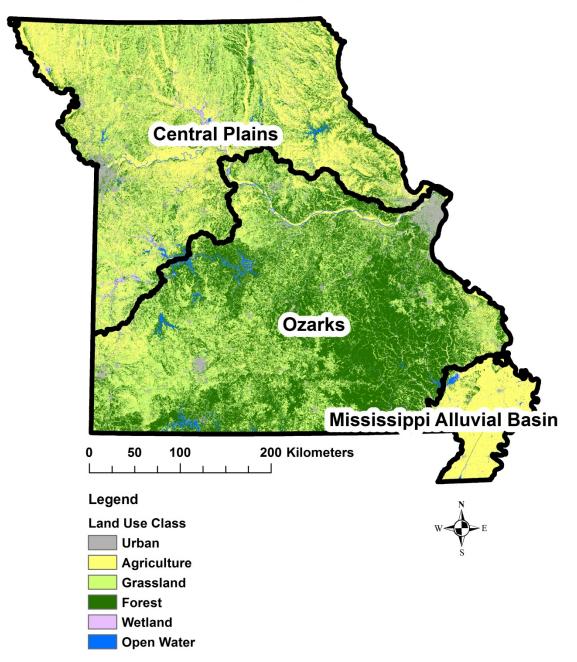
Table 3: Comparison of trait association and species response approaches for classifying species environmental tolerances based on percent agreement of species classifications, odd's ratio and p-value calculated using Fischer's exact test.

| Threat | Percent | Odd's Ratio | P-Value | |
|---------------------|-----------|-------------|---------|--|
| Theat | Agreement | Ouu s ratio | | |
| Habitat Degradation | 65.7 | 3.68 | 0.002 | |
| Temperature | 60.1 | 0.80 | 1.000 | |
| Flow | 46.5 | 0.49 | 0.200 | |

Table 4: Mean number and proportion of vulnerable species at sites within each subregion by index and threat; highest value in bold. P-values tested whether the mean number or proportion of species differed by subregion using a one-way ANOVA. "MAB"=Mississippi Alluvial Basin; "Num."=Number of species; "Prop."=Proportion of species

| Index | Plains | | Ozarks | | MAB | | P-Value | |
|-------------------|--------|-------|--------|-------|------|-------|---------|--------|
| Threat | Num. | Prop. | Num. | Prop. | Num. | Prop. | Num. | Prop. |
| Trait Association | | | | | | | | |
| Habitat | 4.6 | 0.30 | 10.6 | 0.55 | 3.3 | 0.18 | <0.001 | <0.001 |
| Flow | 5.9 | 0.41 | 8.0 | 0.42 | 10.1 | 0.28 | <0.001 | <0.001 |
| Thermal | 3.5 | 0.24 | 8.0 | 0.42 | 5.1 | 0.28 | <0.001 | <0.001 |
| Species Response | | | | | | | | |
| Habitat | 3.0 | 0.19 | 9.2 | 0.51 | 2.4 | 0.16 | <0.001 | <0.001 |
| Flow | 3.5 | 0.25 | 2.3 | 0.13 | 2.1 | 0.14 | <0.001 | <0.001 |
| Thermal | 1.1 | 0.01 | 1.5 | 0.01 | 0.0 | 0.0 | 0.003 | 0.027 |

FIGURES



Missouri Aquatic Subregions and Land Use

Figure 1: Map showing Missouri's aquatic subregions and land use.

Sample Locations

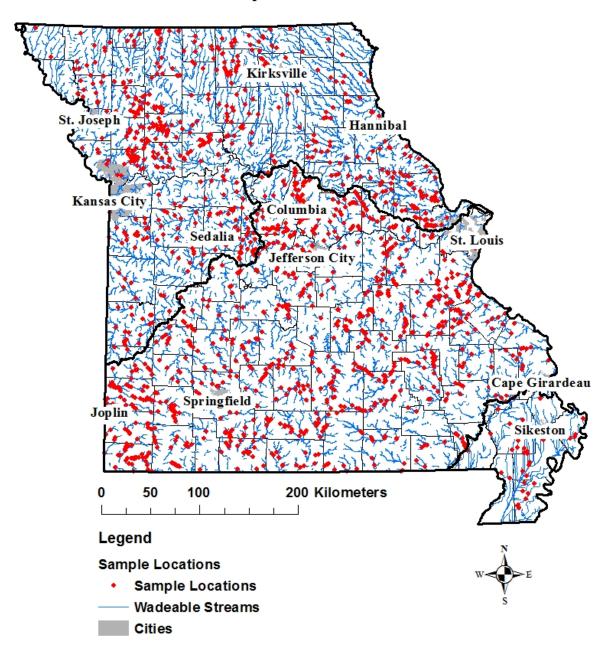


Figure 2: Map of sample locations used in this study for fishes collected from 2000 to 2011 in wadeable streams in Missouri.

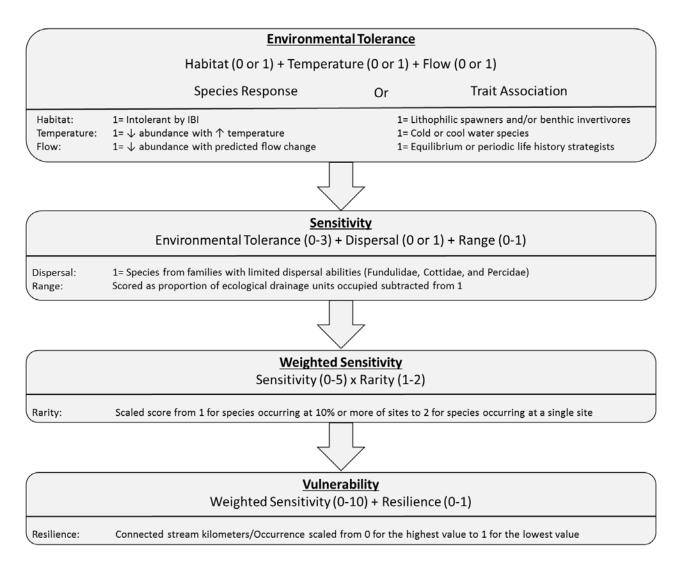


Figure 3: Conceptual diagram of stepwise procedure for calculating species vulnerability scores using the Species Response Index and Trait Association Index. Inputs for dispersal, range, rarity and freshwater resilience were the same for both indices; differences between index calculations occurred only for the environmental tolerance scoring criteria. Numbers in parenthesis indicate potential scoring range. For all "0 or 1" scoring, species meeting listed criteria receive a score of 1, while all others receive a score of 0.

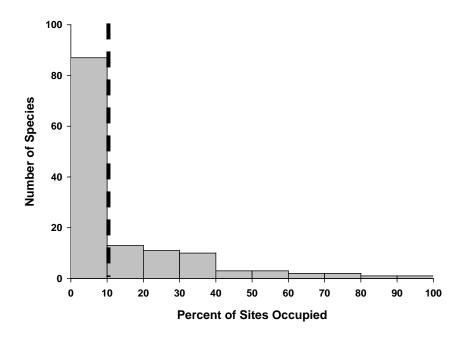


Figure 4: Histogram of species rarity for fish sampled from 1,499 wadeable stream segments in Missouri from 2000 to 2011. The dashed line represents threshold used to classify rare species.

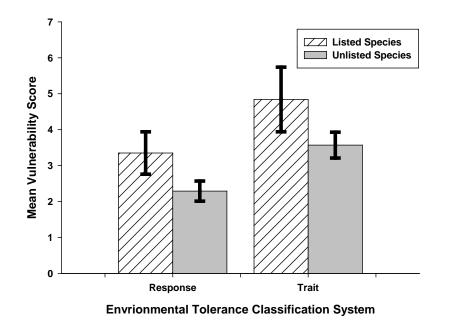


Figure 5: Mean score (+/- 95% Confidence Interval) for listed and unlisted species using the species response and traits association approaches for classifying environmental tolerance.

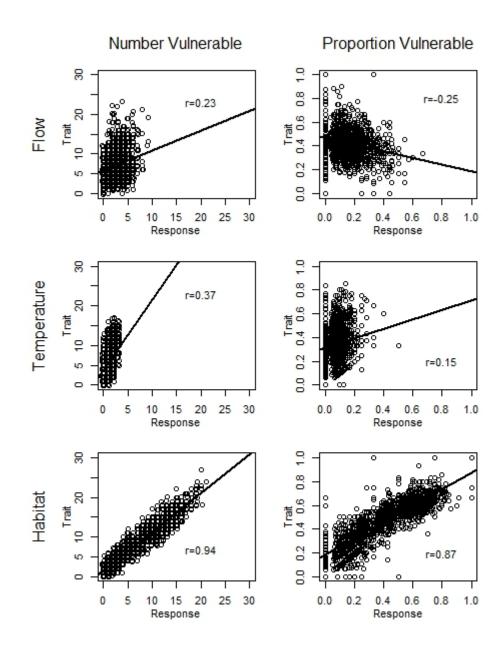


Figure 6: Correlation between the trait and response based approaches for classifying environmental tolerance based on the number and proportion of vulnerable species at each of the study sites, r = Pearson's correlation coefficient.

CHAPTER 2

EVALUATION OF MISSOURI'S CONSERVATION NETWORKS FOR STREAM FISH CONSERVATION

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ABSTRACT

Successful conservation of stream fishes will become increasingly dependent on the use of conservation networks as the effects habitat degradation and climate change increase. We developed a framework for identifying both the most valuable stream segments for fish conservation within established conservation networks, as well as areas outside of conservation networks which best complement what is already being protected. This methodology was applied to three established conservation networks in Missouri, USA: Priority Watersheds, Conservation Opportunity Areas, and the existing conservation network. We also examined patterns in conservation values based on the landscape and habitat characteristics of sites, the impact of constraining prioritization based on the established conservation network compared to using a blank slate approach, how species representation varied in sites selected based on measure of conservation value compared to sites selected based solely on habitat integrity, and how decisions to include other variables (e.g., weighting of species, metric of upstream watershed integrity, and the use of different conservation planning algorithms) may inform conservation planning decisions. There were a number of sites with low scores based on landscape-level habitat integrity, but had conservation values in the top 10%

based on the conservation value approach. Constraining prioritization to established conservation networks generally resulted in lower species representation, however the established networks did include representation of almost all species and provided a more feasible mechanism for implementing large-scale conservation than starting from a blank slate. Species weighting had very little impact on the conservation value of stream segments, likely because highly weighted species generally had smaller distributions and were therefore retained to maintain representation regardless of weight. The inclusion of upstream habitat integrity and the prioritization technique both resulted in differences in the conservation value assigned to sites suggesting that decision makers may need to consider the implications of choices regarding these options on results. We believe this framework can be used in other regions to identify priority sites for the conservation of stream species within and complementary to established conservation networks.

INTRODUCTION

Conservation networks are important for the protection of increasingly imperiled stream communities (Saunders *et al.* 2002; Abell *et al.* 2007; Nel *et al.* 2009a). Habitat degradation, warming stream temperatures, and alterations to the flow regime are expected to contribute to future declines of stream fish (Dudgeon *et al.* 2006). Conservation plans are important for the protection of aquatic biodiversity from these threats, and have been developed for stakeholders around the globe (Abell *et al.* 2007). The primary goal of most aquatic biodiversity conservation plans is to protect the current suite of species or communities, often through reserves or protected areas (Saunders *et al.* 2002; Abell *et al.* 2007). Resources for the establishment and management of reserves and protected areas are often limited; therefore it is important to make scientifically informed decisions to achieve the greatest conservation outcomes.

Effective conservation planning depends on the consideration of both biodiversity and the amount of resources available for protection, management and research to meet goals (Linke *et al.* 2011). Typically, conservation plans for freshwater ecosystems have been centered on the development of a spatial network which supports a targeted set of species or features of interest. Because resources for conservation are often limited, these plans seek to accomplish this while minimizing cost or area by focusing on the complementarity of selected sites (Nel *et al.* 2009a). Recently, there has been a proliferation of research aimed at developing quantitative

methods for creating conservation plans with an emphasis on complementarity (Moilanen *et al.* 2008, 2009a; Hermoso *et al.* 2011; Linke *et al.* 2011). Generally, these tools have been used to design conservation networks or set priorities while treating the project area as a blank slate, that is, areas are prioritized without consideration of any already established conservation networks (Sowa *et al.* 2007; Wenger *et al.* 2009; Strecker *et al.* 2011; Esselman *et al.* 2013; Pool *et al.* 2013). However, in many areas the slate is not blank, as conservation networks have already been established (e.g., state and federal wildlife management areas, refuges, and parks). For many managers and decision makers the flexibility to start from scratch and create new conservation networks or priority areas is not feasible given the costs associated with the creation of new networks, as well as the existing obligations of agencies to its stakeholders (Naughton-Treves *et al.* 2005). In areas where the slate is not blank, evaluation and prioritization techniques which consider established conservation networks constitute a more realistic, practical approach.

This study aims to provide a framework for freshwater conservation planning which incorporates established conservation networks into the planning process, with an application to the state of Missouri. The objectives of this project were to rank wadeable stream segments (based on predicted stream fish representation, species vulnerability to land use and climate change, and the upstream integrity of sites) within and complementary to three established conservation networks in Missouri: the existing conservation network (ECN), Conservation Opportunity Areas (COAs), and Priority Watersheds (PWs). This study also evaluated how decisions related to the inclusion of

species weighting, species specific connectivity requirements, and the method used for prioritizing stream segments affected the results.

The ECN is comprised of public (Missouri Natural Areas, Missouri Department of Conservation (MDC) lands, US Fish and Wildlife Service Refuges, Department of Natural Resources lands, U.S. Forest Service lands, National Park Service lands, and Army Corps of Engineers lands) and private areas (Wetlands Reserve Program lands, The Nature Conservancy Preserves, Missouri Prairie Foundation lands, the Pioneer Forest, and Ozark Regional Land Trust lands), which are managed with a primary purpose of wildlife conservation (Hoskins 2005). In Missouri there are 3,943 ECN units encompassing 13,183 km² and 2,590 km of wadeable streams (Abbitt *et al.* 2004; Figg 2011). COAs consist of 35 units located throughout Missouri that have been identified by MDC as the best locations to achieve all wildlife conservation. These units are a mixture of public and private lands where MDC and 18 partner organizations representing national and state agencies, and national, regional, and local non-profit conservation organizations work together to utilize "technology, expertise and resources for all wildlife conservation" (Missouri Department of Conservation 2012). These COAs encompass 22,855 km² and 4,801 km of wadeable streams (Abbitt *et al.* 2004; Figg 2011). Each COA has a stakeholder team which develops goals and coordinates management and conservation actions for the area and attempts to facilitate conservation actions on both the public and private lands within the areas. PWs were created by the MDC Fisheries Division in 2010 by identifying focal areas across the state for conserving aquatic biodiversity and providing quality areas and opportunities for outdoor recreation. These

areas were selected based on a ranking system which included Aquatic COAs, the presence of streams and impoundments managed by MDC for recreation, local buy-in from landowners, feasibility, and multiple priority achievement in overlapping areas (Corson *et al.* 2010). In total there are 58 Priority Watersheds encompassing 38,931 km² and 8,826 km of wadeable streams (Abbitt *et al.* 2004; Figg 2011).

MATERIALS AND METHODS

Study area

Missouri is comprised of three ecologically unique aquatic subregions: the Central Plains, Ozarks, and Mississippi Alluvial Basin (MAB) (Figure 1; Pflieger 1970; Pflieger and Missouri Deptartment of Conservation 1989; Sowa et al. 2007). Differences in the geology, landform, soils, land cover, and the influence of groundwater between these regions have given rise to unique habitats and fish communities in the streams of each area. The Central Plains encompasses all of northern Missouri and much of the western portion of the state, and characterized by open grassland and agricultural land use (Figure 1; Sowa et al. 2007). The topography is predominately composed of wide, gently sloping valleys (Sowa et al. 2007). Streams are generally characterized as having a relatively low gradient, high turbidity, and fine silt and sand substrates (Sowa et al. 2007). Most of the southern portion of Missouri is considered the Ozarks, which is predominately composed of a mix of forested areas and open pastureland (Figure 1; Sowa *et al.* 2007). The topography has high relief and rugged terrain. Streams in the Ozarks tend to have higher average stream gradients with gravel, cobble, or bedrock as dominant channel substrates with sand and silt substrates in slower moving areas and

pools (Sowa *et al.* 2007). The far southeastern portion of the state is the Mississippi Alluvial Basin (MAB), which has low relief and low gradient streams (Sowa *et al.* 2007). Most of this region has been channelized and ditched for agricultural production (Figure 1). The substrate varies throughout the subregion but generally the larger and slower moving streams are characterized by fine silt while streams with faster flow are comprised of sand and small gravel substrates (Sowa *et al.* 2007).

Data

Fish community data was provided by MDC and included records from 1990 to 2011 (Figure 2). Data collected between 1990 and 1999 was contained in the MDC Fish Community Database, while data collected between 2000 and 2011 was contained in the Resource Assessment and Monitoring Program (RAM) database. Data used for this study was limited to fish community collections from wadeable streams (2nd-5th order and classified as permanent), and had at least 0.5 hours of effort using seines and/or electrofishing. This allowed for a scope of inference which included all wadeable, permanent streams in Missouri (Figure 2). All 769 samples within the MDC Fish Community Database were representative of community sampling efforts; while all 1,107 samples within the RAM database were sampled using a standard procedures which used block nets to enclose the sample area which was then thoroughly sampled via electrofishing and seining (Fischer and Combes 2003). Sites were selected for RAM sampling based on a stratified random approach which selected wadeable stream segments throughout Missouri (Fischer and Combes 2003). In this study both the RAM and Fish Community Databases were used to account for stream fish occurrence in a

presence/absence format. In addition to the above databases that contained information on community samples, point data obtained from the Missouri Natural Heritage Program database, which was collected between 1990 and 2011, was used for species for which models could not be developed.

Environmental data was provided by the Missouri Resource Assessment Partnership (MoRAP). This geospatial data was linked to each confluence to confluence stream segment and included 27 variables related to biogeography, stream features, local landscape, upstream landscape, and anthropogenic impacts (Appendix 5; Abbitt *et al.* 2004). These variables were selected because they have been linked to fish species distributions in Missouri and elsewhere (Sowa *et al.* 2007; Strecker *et al.* 2011).

Species distribution models

Species distribution models were developed to determine representation of each species in each wadeable stream segment in Missouri. Models were developed using an ensemble approach that averaged the results of four component models which met minimum evaluation standards. The four component models included multivariate adaptive regression splines (MARs), generalized additive models (GAMs), boosted regression trees (BRTs), and random forest models. Species models were developed separately for each of Missouri's aquatic subregions because species exhibit different relationships with landscape features between subregions (Sowa *et al.* 2007). Species presence or absence served as the response variable while predictor variables were drawn from both continuous and categorical environmental data (Appendix 5). Models

were developed for species which had at least 40 occurrences within a subregion. All models were developed using R statistical software using the 'earth' package (MARs; Milborrow 2014), 'gam' package (GAM; Hastie 2014), 'gbm' package (BRT; Ridgeway 2013), and the 'randomForest' package (random forest; Liaw *et al.* 2014). Each model was built from a random subset of 70% of the species occurrence data, with the remaining 30% being used for model evaluation.

Model accuracy was evaluated using the area under the receiver operator characteristic curve (AUC), calculated using the 'ROCR' package in r (Sing *et al.* 2013), model bias (number of occurrences predicted compared to observed expressed as a percentage), and model fit as the mean absolute error (MAE) of a calibration curve with 10% probability of occurrence bins. Ensemble models, which predicted probability of occurrence for all species in each wadeable stream segment in Missouri, were created by averaging the predictions of all component models which met each of the minimum evaluation standards: AUC \geq 0.6, bias of +/- 25%, and MAE of \leq 0.125.

Species representation

Species representation within Missouri's established conservation networks was assessed by determining the number of stream segments a species was predicted to occupy within each conservation network. Species occupancy was accounted for in two ways. For species which met the modeling requirements predicted probabilities of occurrence were drawn from the species distribution models. For species, for which acceptable models were not created, observed occurrence records were used to assign

stream segments a probability of occurrence of 1 where the species had been collected, and the species frequency of occurrence for each ecological drainage unit (EDU) was then used to assign probabilities of occurrence to all unsampled stream segments. This was accomplished for each of the 17 ecological drainage units (EDUs) in Missouri by dividing the number of occurrences from the number of samples in each EDU. Predicted number of occupied stream segments within each of the established conservation networks was calculated for each species by summing the probabilities of occurrence for all stream segments which intersected a conservation network.

Conservation value

Stream segments were prioritized within and complementary to Missouri's conservation networks based on a measure of conservation value. Conservation value rankings were calculated based on weighted species representation while accounting for upstream watershed integrity using Zonation conservation planning software version 3.1 (Moilanen *et al.* 2012). Conservation values were calculated via two zonation algorithms which calculated the proportional loss of species representation based on the removal of each stream segment. The stream segment with the lowest conservation value was removed, and conservation values for all remaining cells were recalculated. This process is done iteratively until only a single stream segment remains producing a hierarchy of conservation values from the least to most valuable stream segment. This analysis was completed for a number of scenarios which incorporated different methods of accounting for representation, weighting species, and upstream

integrity for each of the three conservation networks as well as the blank slate approach (i.e., assumes no established conservation networks).

Species representation was accounted for using two distinct methods: core-area Zonation and with the additive benefit function. Core-area Zonation by removing the cell which minimizes biological loss by selecting the stream segment with the lowest value for the most valuable species occurrence in that cell; this approach emphasizes unique species occurrences and generally results in better representations of uncommon species (Moilanen *et al.* 2012). The additive benefit function minimizes biological loss by selecting the stream segment that results in the lowest marginal loss of all species in a given cell for removal (Moilanen *et al.* 2012). This method emphasizes species richness and generally results in higher average species representation but lower levels of representation for uncommon species (Moilanen *et al.* 2012).

Species representation was weighted in three different ways to capture a number of options by which decision makers may value the representation of species. These weights were based on species vulnerability, species listing status, or all species weighted equally. Species were weighted based on vulnerability and listing status to emphasize the selection of areas for species in greatest need of conservation, and was compared to the analysis where species had equal weights. Vulnerability weight was based on a species vulnerability to habitat degradation, warming stream temperatures, and alterations to the flow regime while also considering a species dispersal ability, range size, rarity, and range-wide fragmentation (Chapter 1). The trait association index

developed in Chapter 1 was used to calculate our vulnerability scores because it could assess all species of interest (Chapter 1). Vulnerability weights were scaled from 1 (lowest vulnerability score) to 2 (highest vulnerability score). Conservation prioritization was also completed using state listing status as a weight, where species of conservation concern received weights of 2, and unlisted species received weights of 1, as well as with all species weighted equally.

Species-specific penalty curves were developed, adapting the procedure developed by Moilanen et al. (2008), to account for the potential effect of habitat degradation of upstream sites on the value of a local stream segments fish community (Figure 3). Species-specific penalty curves were developed using the RAM fish community data by dividing observed species occurrences into three classes based on upstream habitat degradation (Percent of upstream watershed area classified as urban or agricultural land use). These classes were low (0-33%) which served as a baseline, medium (>33-66%), and high (>66-100%). Frequency of occurrence within each class was calculated for each species which had a minimum of ten occurrences, were then divided by the baseline occurrence rate, and values were then rounded to the nearest value of 1, 0.66, or 0.33 to represent the multiplicative change in biological value of a stream segment based on the remaining proportion of its upstream watershed. Values for each class were restricted to being no higher than the previous, more intact class, when this exception occurred the more intact class was assigned the value of the less intact class. This was done to provide a more conservative estimate of the loss of biological value and to eliminate situations in which a loss of upstream sites would

result in an increase in biological value. This exception occurred for 20 of the 118 species which were evaluated. Fifteen species did not have a sufficient number of occurrences (10) for analysis so were analyzed without accounting for upstream habitat integrity. Integrity curves were then created with x-axis values of 0 (low class), 0.5 (medium class), and 1 (high class) with the associated y-axis values representing multiplicative changes in value (1, 0.66, 0.33).

Stream segments were ranked both within and complementary to Missouri's established conservation networks, which forced stream segments within each conservation network to be retained until all areas outside of the network were removed (Moilanen et al. 2012). This process created a ranking of stream segments outside of the established conservation network from least to most valuable respective of what is represented within the established conservation networks. Stream segments outside of the conservation network with predicted occurrences of species not well represented within the conservation network received high conservation values, particularly for species with high weights, whereas, sites with species which are already well represented within the conservation network received lower values. Stream segments within the conservation network were then removed, creating a ranking within each established conservation network. This analysis was conducted separately for Priority Watersheds, Conservation Opportunity Areas, and the existing conservation network in Missouri. In addition, this analysis was completed without accounting for any established conservation networks to develop a blank slate solution for the purpose of comparison. This analysis was conducted for each of the established conservation

networks, and the blank slate, using all possible combinations of accounting for species representation (core-area and additive benefit), species weighting (vulnerability, listing status, all equal), and upstream watershed integrity (with and without).

This study analyzed whether areas of high conservation value differed by landscape-level habitat features including aquatic subregion, ecological drainage unit, Strahler stream order, segment gradient, and percent of watershed in forest, agriculture, grassland, and urban land cover for each of the three aquatic subregions, as land use was substantially different among subregions (Figure 1). Continuous features were broken down into two classes; top 10 percent of sites and remaining 90 percent of sites, while categorical features consisted of classes based on their discrete features. A one way analysis of variance (ANOVA) was performed for each feature to determine whether mean conservation values differed by segment characteristics.

The representation of species in areas selected through the use of the most basic prioritization analysis (core area, equal weights, no upstream integrity consideration) was compared to the selection of the least degraded stream segments to determine the potential benefits of applying a systematic, representation based approach. The level of degradation of a stream segment was classified based on a habitat threat index (HTI) for Missouri and surrounding states (Annis *et al.* 2009). Within each of the three established conservation networks the top ten percent of stream segments based on greatest conservation value and the top ten percent based on lowest HTI score (i.e., lowest landscape-level threats) were selected for each of the three subregions. The

number of species predicted to be lacking representation in each solution, the percent of species predicted to occupy the greater number of stream segments in the solution derived using conservation value versus HTI scores, and the mean, maximum, and minimum percent difference between the conservation value and HTI derived solutions were all calculated. The top 2.5% of complementary areas selected based on the two approaches were analyzed for each established network in each of the aquatic subregions by calculating the percent of species with more predicted occupied stream segments in the solution derived using conservation value, and the average, maximum, and minimum percent differences from the HTI derived solution to the conservation value derived solution.

The level of species representation achieved by the established conservation networks was compared to an optimal network generated using the blank slate approach. The number of species expected to be represented within each established conservation network and the corresponding optimal networks were calculated. This study also determined the proportion of each species predicted occurrences within each conservation network as well as within an optimal conservation network of the same size, which was used to identify the species with the lowest proportion of its occurrences protected within each of the established conservation networks, as well as the optimal networks of the same size. The mean proportion of species occurrences protected within a conservation network was also calculated for each of the established networks and the corresponding optimal networks. These results were grouped by network and conservation prioritization technique (Cell removal algorithm: core area or

additive benefit) and were analyzed using a three way ANOVA to determine if there was a difference in both the minimum and mean proportions of species occurrences between established networks and the optimal network while including connectivity and species weighting scheme as factors.

A sensitivity analysis was conducted to determine the effects of the prioritization technique, inclusion of upstream connectivity, and species weighting, on conservation value scores. Percent congruence for sites ranked in the top 1%, 5%, as well as each 10% increment between 0 and 100% of the prioritization results were calculated within and complementary to each of the conservation networks . These comparisons were made at a pairwise level in which all settings and inputs were the same except the input or setting analyzed, which allowed us to determine how sensitive the prioritization analysis is to the use of different prioritization algorithms, inclusion of connectivity, and the weighting of species.

RESULTS

Species distribution models were developed for 40 species in the Plains subregion, and 68 in the Ozarks subregion. All species for which attempts were made to create models, except Black Bullhead *Ameiurus melas* and Green Sunfish *Lepomis cyanellus* (both Plains models), had at least one of the four ensemble models which met minimum evaluation standards and were therefore represented via a distribution model (Appendix 6). All remaining species representations (44, 64, and 50 species for the Plains, Ozarks, and Mississippi Alluvial Basin respectively) had less than 40 occurrences

within a subregion and were therefore classified based on occurrence locations and frequency of occurrence data. Based on the three model evaluation criteria (AUC, BIAS, and MAE) Random Forest models generated the most acceptable component models (95), followed by BRTs (89), GAMs (81), and finally MARS (76) (Table 1). The ensemble approach allowed 108 species distribution models to be created, which was a 42 percent increase over MARS alone, 33 percent increase over GAMs alone, 21 percent increase over BRTs alone, and 14 percent increase over Random Forest models alone. More detailed information regarding the species distribution models can be found in the appendices including: regional species model list (Appendix 6), ensemble components (Appendix 7), component model evaluation statistics (Appendix 8), and maps and shapefiles of results (Appendix 9).

All species were predicted to occur in at least 1 stream segment within each of Missouri's established conservation networks (Appendix 10). In stream segments within the established conservation networks 6, 8, and 7 species were predicted to be found in less than 10 stream segments for PWs, the ECN, and COAs respectively. There were 97, 90, and 90 species which were predicted to occupy greater than 100 stream segments in PWs, the ECN, and COAs respectively.

Conservation value was variable but showed patterns based on subregion and EDU (Ps<0.01; Appendix 11). All three subregions contain stream segments with both high and low conservation values (Figure 4; Appendix 11). The MAB had the highest mean conservation value (0.79) followed by the Ozarks (0.64) and Plains (0.30)

(P<0.001). In the Plains average conservation value scores ranged from 0.42 in the Nishnabotna/Platte EDU to 0.16 in the Osage/South Grand EDU (P<0.001). In the Ozarks average conservation values ranged from 0.89 EDU in the Neosho EDU to 0.41 in the Moreau/Loutre EDU (P<0.001). Finally, in the Mississippi Alluvial Basin average conservation values ranged from 0.96 in the Black/Cache EDU to 0.78 in the St. Francis/Little EDU (P<0.001). Average conservation value tended to increase with increasing Strahler order for the Plains (0.23 for 2nd order to 0.44 for 5th order; P<0.001) and the Ozarks (0.56 2nd order to 0.76 for 5th order; P<0.001), and the MAB (0.78 for 5th order to 0.80 for 2nd order; P<0.001), but the differences in the MAB were likely not ecologically significant. The top 10 percent of highest conservation value sites in the Plains had higher forest cover in the watershed (15.6% to 12.9%; P<0.001), lower agriculture (39.2% to 43.7%; P<0.001), and lower grassland (39.2% to 43.7%; P<0.001) than sites in the lower 90% of conservation values. In the more forested and less agriculturally-dominated Ozarks the top 10 percent of highest conservation value sites had lower levels of agriculture in the watershed (2.1% to 4.9%; P<0.001), and lower stream segment gradients (2.3 m/km to 3.1m/km; P<0.001), but forest cover was similar (48.3% to 49.7%; P<0.001). In the Mississippi Alluvial Basin high value stream segments had more forest in their watersheds (13.3% to 6.8%) and less agriculture (72.7% to 80.4%) in this agriculturally-dominated subregion. These results suggest that land use and land cover was related to conservation value, but was region-specific.

Stream segments selected based on conservation value resulted in greater species representation, particularly for uncommon species when compared to sites

selected based on habitat integrity alone (HTI scores) (Table 2). The number of species lacking representation in the top ten percent of sites within established conservation networks based on conservation value were one species in Priority Watersheds, two in the existing conservation network, and five in Conservation Opportunity Areas, while the number of species lacking representation in a corresponding number of sites selected based on HTI were 10 in Priority Watersheds, 13 in the existing conservation network, and 17 in Conservation Opportunity Areas. Between 70 and 77 percent of species were predicted to have higher levels of representation in the top ten percent of segments within conservation networks selected using the conservation value framework than those selected based on habitat integrity alone (Table 2). Additionally, the average the use of conservation value retained 166% more occurrences per species than using HTI for Priority Watersheds, 311% more for Conservation Opportunity Areas, and 190% more for the existing conservation network. Uncommon species had the largest increases in number of occurrences based on the selection of sites using conservation value as opposed to using habitat integrity with increases of 6,145% (0.2 predicted occurrences for sites selected based on HTI vs 15 predicted occurrences for sites selected based on conservation value) for the Redspot Chub *Nocomis asper* in the existing conservation network, and 7,099% (0.4 vs 28.9 occurrences) and 16,122% (0.1 vs 21.8 occurrences) for the Niangua Darter *Etheostoma nianguae* in Priority Watersheds and Conservation Opportunity Areas respectively. Several common species were less well represented in sites selected based on conservation value rather than HTI score including the Mottled Sculpin Cottus bairdii in Priority Watersheds (65% less

predicted occurrences; 120 vs 47 sites), the Creek Chubsucker *Erimyzon oblongus* in Conservation Opportunity Areas (61%; 55 vs 21) and the existing conservation network (65%; 55 vs 19). The amount of species which were had more occurrences when selections were made using conservation value ranged from 70% in Conservation Opportunity Areas to 73% in both Priority Watersheds and the existing conservation network. The average percent difference ranged from 11% in Priority Watersheds to 14% in Conservation Opportunity Areas, with maximum differences ranging from 127% in Conservation Opportunity Areas to 321% in the existing conservation network and minimum differences ranging from -8% for the existing conservation network to -16% for Conservation Opportunity Areas.

Established networks retained 4.2 to 10.0% higher proportion of representation than the blank slate alternative depending on network for the species which lost the greatest proportion of their original representation in each solution developed using the additive benefit function (Table 3). When core-area Zonation was used to create blank slate alternatives there was no significant difference for Priority Watersheds (-1.0%; P=0.195), but both Conservation Opportunity Areas (-1.2%), and the established conservation network (-3.3%) had decreased representation when compared to the blank slate alternatives (Table 3; Ps<0.001). The average proportion of representation lost across all species was less in the optimal network alternatives than in the established networks using both additive benefit (Table 3; -27.4%, -11.7%, and -23.5% change for Priority Watersheds, Conservation Opportunity Areas, and the existing conservation network respectively), and core-area Zonation (Table 3; -15.9%, -5.7%, and

-12.6% change for Priority Watersheds, Conservation Opportunity Areas, and the existing conservation network respectively). Therefore, developing a network using a blank slate approach typically resulted in a lower reduction of species distribution compared to established conservation networks.

Our assessment of the influence of connectivity, prioritization technique, and species weighting on the conservation ranking of stream segments revealed that species weighting and connectivity had less influence on the results than prioritization technique. Conservation scores of stream segments determined with and without connectivity revealed high levels of congruence even among the most valuable sites (Figure 5A). Congruence analysis of the conservation scores of stream segments using core-area Zonation compared to the additive benefit function revealed lower levels of congruence suggesting the choice of prioritization technique can have substantial effects on the outcome (Figure 5B). Finally, congruence of conservation rankings was relatively high between all pairwise combinations of species weighting (none, vulnerability, listing status) suggesting that the weighting of species had only a moderate effect on the conservation solutions (Figure 5C, D, E).

DISCUSSION

This study provides a framework that may be a more realistic and effective approach for prioritizing stream fish conservation efforts over a broad spatial scale when conservation networks have already been established. This framework, may serve as a more feasible option for conservation planning than previous work in the area

which has largely used a blank slate approach (Sowa et al. 2007; Wenger et al. 2009; Strecker et al. 2011; Esselman et al. 2013; Pool et al. 2013). Budgets for agencies tasked with managing and creating conservation areas and protecting stream fish are often limited, necessitating the use of efficient prioritization schemes for achieving the greatest conservation outcomes (Naughton-Treves *et al.* 2005). Therefore, the ability to prioritize stream fish conservation within established networks may be critical, as agencies typically do not have the resources necessary to implement stream fish conservation actions for all stream segments within conservation networks. By incorporating established conservation networks in the planning process, stream segments which complement what is already being protected can also be identified. This method emphasizes the selection of areas which have occurrences of species which may not already be well represented within an established network resulting in an expanded network with a more comprehensive and diverse assemblage of species. This framework allows managers and decision makers to make more informed decisions regarding the spatial prioritization of stream fish conservation.

Analysis revealed that Missouri's established conservation networks typically do contain the vast majority of fish species in the state. All 133 species which were evaluated were predicted to be represented within each of Missouri's established conservation networks except Black Buffalo, *Ictiobus niger*, a species which is more commonly found in rivers than the wadeable streams that were sampled for this study (Pflieger 1997). Protecting multiple areas to allow for redundancy in case of catastrophic declines and changes in habitat due to climate change, invasive species, or

anthropogenic disturbances may be needed to ensure species persistency (Stein et al. 2000), and our study revealed that the majority (68-72%, depending on conservation network) of species were predicted to occur in over 100 stream segments in each of the established conservation networks suggesting they are relatively secure. However 5 to 7 species (depending on conservation network) were predicted to occur within less than 10 stream segments, suggesting their long-term protection is less secure. These less secure species often had restricted distributions such as the Channel Darter Percina copelandi, Arkansas Saddled Darter Etheostoma euzonum, Flier Centrarchus macropterus and Bluntface Shiner Cyprinella Camura, were rare or difficult to detect such as the Southern Brook Lamprey *Icthyomyzon gagei*, and Northern Brook Lamprey *Icthyomyzon fossor*, or were primarily found in riverine habitats rather than wadeable streams such as the Black Buffalo Ictiobus niger, Western Silvery Minnow Hybognathus argyritis, Plains Minnow Hybognathus placitus, and Highfin Carpsucker Carpiodes velifer (Pflieger 1997; Galat et al. 2005). When possibilities for network expansion arise targeting areas with known or predicted occurrences of underrepresented species may improve the long-term outlook for those species, and bolster the comprehensiveness of the established network.

The use of surrogates, such as landscape and habitat characteristics, is widely used in conservation planning, particularly when access to biological data is limited (Nel *et al.* 2009b). The findings of this study suggest that results from habitat-based assessment may not produce the same results as assessments based on species representation. High value areas which are excluded when sites are selected based on

landscape and habitat surrogates are likely to include species which are restricted to non-surrogate habitat classes and rare species. This study found that stream segments of both high and low conservation value were distributed throughout the state and in a variety of stream sizes, ecological subregions, drainage basins, and land use classes. Selecting sites based only on high value habitat surrogates would have meant that some stream segments in the top 10 percent of conservation values, belonging to groups associated with the lowest average conservation values (Plains aquatic subregion, the Osage/South Grand, Grand Chariton, and Blackwater/Lamine drainages, 2nd order streams, and agricultural watersheds) would have been excluded. Species restricted to these areas, and rare species would have much lower representations if a prioritization system using only landscape and habitat surrogates was used. Although the use of landscape and habitat surrogates are a useful alternative when sufficient data on species representation is not available (Trakhtenbrot and Kadmon 2005), a systematic conservation planning approach which includes species representation may be important for comprehensive species conservation (Margules and Pressey 2000; Linke et al. 2011). In cases where species representation data is not available the results of this study suggest that landscape and habitat based surrogates do a reasonable job of identifying sites with high conservation values. However, when sufficient biological data is available, representation based, systematic conservation planning is likely to result in solutions which better represent rare species and species which occupy habitats not associated with high value surrogates.

One commonly used method for creating protected areas is the selection of areas which are pristine or have minimal anthropogenic impacts, often because these areas are rugged, isolated, or have limited economic value (Margules and Pressey 2000). A number of studies have developed tools for assessing the ecological integrity of catchments (Mattson and Angermeier 2007; Annis et al. 2009; Paukert et al. 2011), and although we believe this information is a valuable component of conservation planning, our results suggest that relying on threat data alone, without consideration of species representation, may result in limited or no protection for many species. Using this studies measure of conservation value for prioritizing sites yielded much higher predicted levels of species representation (9 to 12 more species were represented in the top 10% of sites within each conservation network) than if stream segments would have been selected based solely on a measure of habitat integrity (Annis et al. 2009). This suggests that systematic, representation based conservation planning is likely to maintain higher levels of representation for the majority of species (species averaged between 166-311% greater representation in the top 10% of sites within each of conservation network) and particularly so for uncommon species both within and complementary to established conservation networks. In order to maintain comprehensive species representation prioritization of sites based on threat data alone is likely insufficient.

Systematic conservation planning often considers a variety of variables, in addition to species representation, to account for factors which influence conservation success (Moilanen *et al.* 2008; Leathwick *et al.* 2010; Strecker *et al.* 2011; Pool *et al.*

2013). The weighting of species is commonly used in conservation planning which allows emphasis to be placed on sites which are important for priority species (Moilanen *et al.* 2005; Early and Thomas 2007; Gordon *et al.* 2009; Carroll *et al.* 2010; Summers *et al.* 2012). However, the results showed that weighting species had a relatively minor effect on the conservation values of stream segments. Species which were weighted highly also tended to be the least common, and because this study's methodology placed emphasis on maintaining representations of all species, stream segments with occurrences of uncommon species were given high conservation priority, regardless of weighting. The lack of sensitivity to species weighting suggests that uncertainty or differences in weighting schemes are unlikely to have substantial impacts on conservation priorities for others employing this framework.

Upstream habitat integrity appeared to have an intermediate impact on the prioritization of sites both within and complementary to established conservation networks. Stream fish communities are heavily influenced by conditions in the upstream watershed (Weaver and Garman 1994; Wang *et al.* 1997, 2001; Wenger *et al.* 2008). Many conservation plans incorporate upstream habitat integrity requirements (Moilanen *et al.* 2008; Hermoso *et al.* 2011; Strecker *et al.* 2011; Esselman *et al.* 2013), but there has been little evaluation of how the inclusion of connectivity influences results. The inclusion of connectivity in these analyses tended to increase the value of upstream segments and showed some additional clustering when compared to results where connectivity was not included.

The results of this study suggest that the choice of prioritization technique had a substantial impact on the results of the analysis. A variety of prioritization techniques have been developed which assign conservation values in a number of unique ways (Moilanen 2007; Moilanen et al. 2009b; Watts et al. 2009). The use of two prioritization techniques, which 1) emphasizes unique species representation (core area), and 2) emphasizes species richness (additive benefit; Moilanen et al. 2012) showed substantial differences (<60% congruence for top 10 percent of sites), and the proportion of representation lost by the average species and the species which lost the greatest proportion of its predicted occurrences. Prioritizations performed using the core-area implementation had higher levels of representation for the worst off species, while those performed using the additive benefit function maintained higher levels of representation for the average species, as was expected based on the literature (Moilanen et al. 2012). The sensitivity of conservation value rankings to the choice of prioritization technique suggests that researchers and decision makers should carefully consider their objectives when determining which approach to use. If the primary objective of the prioritization is to maintain representation of all species the core-area implementation is likely the most appropriate approach, while if the primary objective is to select stream segments with higher species richness and retain higher average levels of species representation implementation the additive benefit function may be a better approach.

Our study developed tools which can be used by managers and decision makers to prioritize management, land acquisitions, and guide the formation of partnerships for

stream fish conservation in Missouri, and an approach for conservation prioritization which can be applied to other regions. Stream fish face a number of threats including habitat degradation, climate change, invasive species, and habitat fragmentation (Strayer and Dudgeon 2010). In order to conserve aquatic biodiversity in the face of these threats managers undertake a variety of management actions such as riparian restoration, watershed management, and barrier removal (Roni *et al.* 2002). Conservation planning techniques can help identify where these actions can be taken in order to achieve the greatest returns. The conservation value of complementary sites to established conservation networks may be used to identify opportunities for land acquisition or partnerships which offer the greatest value.

We believe this framework for incorporating established conservation networks into the conservation prioritization process can be used for aquatic biodiversity conservation for regional or national prioritization projects around the globe. In the United States data on protected areas is widely available through the National Gap Analysis Program Protected Area Database (DellaSala *et al.* 2001), which contains 635,500 geospatial records from 49 agency types including state and federal agencies, municipal governments, and non-profit organizations for all 50 states (Duarte 2012). Protected and priority areas, such as South Africa's National Freshwater Ecosystem Priority Areas (Nel *et al.* 2011), Europe's Natura 2000 sites (Commission of the European Communities 2002), and State Wildlife Action Plans in the United States (Pugh and Hall 2006) among many others represent a vast set of established networks upon which this framework for prioritizing stream fish conservation could be applied. The ability to

consider established conservation networks during the prioritization process adds realism to the systematic conservation planning process. In regions where sufficient data exists to estimate the distributions of aquatic biodiversity, this framework can be utilized to inform management decisions within established networks, and can guide land acquisition and partnerships in selecting areas which best complement what is already being protected.

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TABLES

Table 5: Summary statistics for the evaluation metrics for each of the four component model types. Accept= Percent of models which met minimum evaluation criteria; AUC= Area under the receiver operating characteristic curve; MAE= Mean absolute error; MARS= Multivariate adaptive regression splines; GAM= Generalized additive model; BRT= Boosted regression tree; RndmFrst= Random forest

| | AUC | | Bias | | MAE | | |
|----------|-------------|----------|-------------|--------|-------------|--------|--|
| Model | Mean | Accept | Mean | Accept | Mean | Accept | |
| | (Range) | / locept | (Range) | neeepe | (Range) | | |
| MARS | 0.78 | 92.6 % | 1.00 | 83.3 % | 0.08 | 88.0% | |
| | (0.44-1.00) | | (0.52-1.73) | | (0.03-0.20) | | |
| GAM | 0.80 | 94.4 % | 0.97 | 84.3 % | 0.08 | 91.7 % | |
| | (0.52-0.97) | | (0.57-1.85) | | (0.02-0.21) | | |
| BRT | 0.83 | 99.1 % | 1.05 | 87.9 % | 0.07 | 95.3 % | |
| | (0.48-1.00) | | (0.74-2.49) | | (0.03-0.17) | | |
| RndmFrst | 0.83 | 97.2 % | 0.99 | 89.8 % | 0.08 | 94.4 % | |
| | (0.53-1.00) | | (0.61-1.59) | | (0.04-0.24) | | |

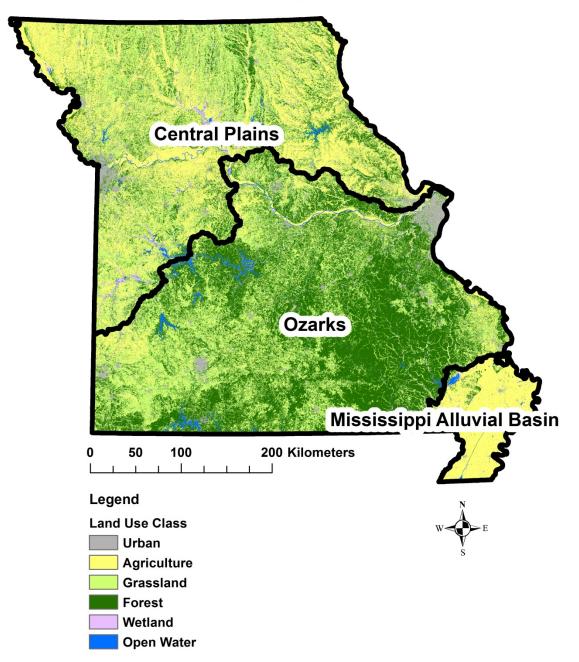
Table 6: Comparison of sites selected based on habitat threat index (HTI; Annis *et al.* 2009) versus conservation value scores. Percent difference was calculated based on the number of stream segments predicted to be occupied for a species for segments selected based on conservation value versus segments selected based on HTI, where positive % differences represent increased representation when using conservation value and negative % differences represent decreased representation when using conservation value. The number of species unrepresented is the number of species which are not predicted to be represented using either HTI or conservation value for stream segment selection. PW=Priority Watersheds; COA=Conservation Opportunity Areas; ECN=existing conservation network

| | PW | COA | ECN | | |
|--|-------|--------|-------|--|--|
| Top 10% of Conservation Network | | | | | |
| % Species Better Represented by Conservation Value | 77 | 73 | 70 | | |
| Mean % Difference | 166 | 311 | 190 | | |
| Max % Difference | 7,099 | 16,122 | 6,145 | | |
| Min % Difference | -61 | -61 | -65 | | |
| # Species Unrepresented (HTI) | 10 | 17 | 13 | | |
| # Species Unrepresented (Conservation Value) | 1 | 5 | 2 | | |
| Entire Conservation Network and Top 2.5% Complementary Areas | | | | | |
| % Species Better Represented by Conservation Value | 73 | 70 | 73 | | |
| Mean % Difference | 11 | 14 | 11 | | |
| Max % Difference | 199 | 127 | 321 | | |
| Min % Difference | -14 | -16 | -8 | | |

Table 7: Comparison of the mean proportion of distribution lost for 1) the species which has lost the highest proportion of distribution and 2) the average proportion of distribution lost when restricted to each of the conservation networks compared to proportional distribution lost when restricted to an optimal network of equivalent size. P values calculated using a three way anova based on weighting method, connectivity, and current/optimal network. PW=Priority Watershed, COA=Conservation Opportunity Area, ECN=Existing Conservation Network. Method acronyms: CAZ=Core Area Zonation, AdBen=Additive Benefit Function.

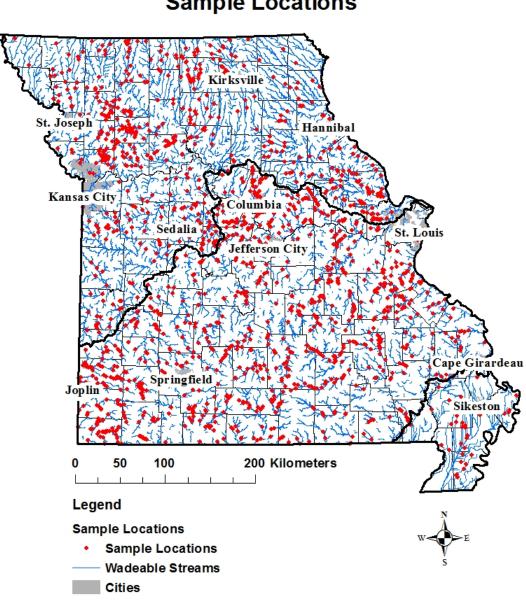
| | | Species with Highest Proportion Lost | | | Average Proportion Lost | | | | |
|---------|--------|--------------------------------------|---------|---------|-------------------------|---------|---------|---------|---------|
| | | Network | Optimal | Percent | | Network | Optimal | Percent | |
| Network | Method | Mean | Mean | Change | Ρ | Mean | Mean | Change | Ρ |
| PW | CAZ | 0.899 | 0.890 | -1.0% | 0.195 | 0.730 | 0.614 | -15.9% | <0.001 |
| | AdBen | 0.899 | 0.991 | 10.2% | < 0.001 | 0.730 | 0.530 | -27.4% | < 0.001 |
| COA | CAZ | 0.955 | 0.944 | -1.2% | 0.013 | 0.804 | 0.758 | -5.7% | <0.001 |
| | AdBen | 0.955 | 0.995 | 4.2% | < 0.001 | 0.804 | 0.710 | -11.7% | < 0.001 |
| ECN | CAZ | 0.932 | 0.901 | -3.3% | 0.006 | 0.744 | 0.650 | -12.6% | < 0.001 |
| | AdBen | 0.932 | 0.992 | 6.4% | < 0.001 | 0.744 | 0.569 | -23.5% | <0.001 |

FIGURES



Missouri Aquatic Subregions and Land Use

Figure 7: Map showing Missouri's aquatic subregions and land use classifications.



Sample Locations

Figure 8: Map of fish community sample locations, collected in Missouri from 1990 to 2010, which were used in this study.

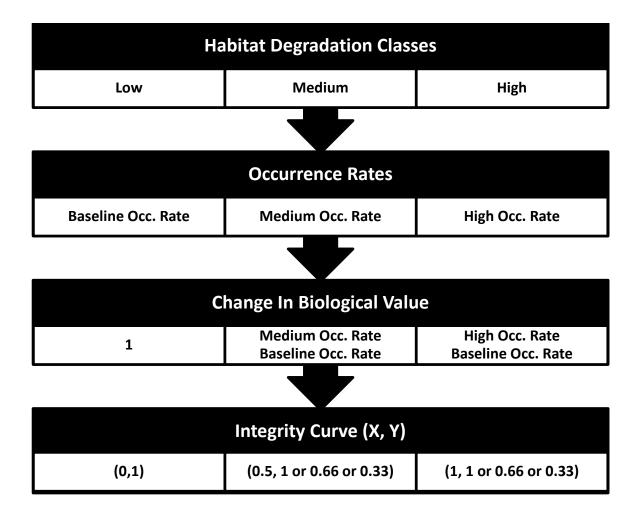


Figure 9: Framework used for developing upstream habitat integrity penalty curves.

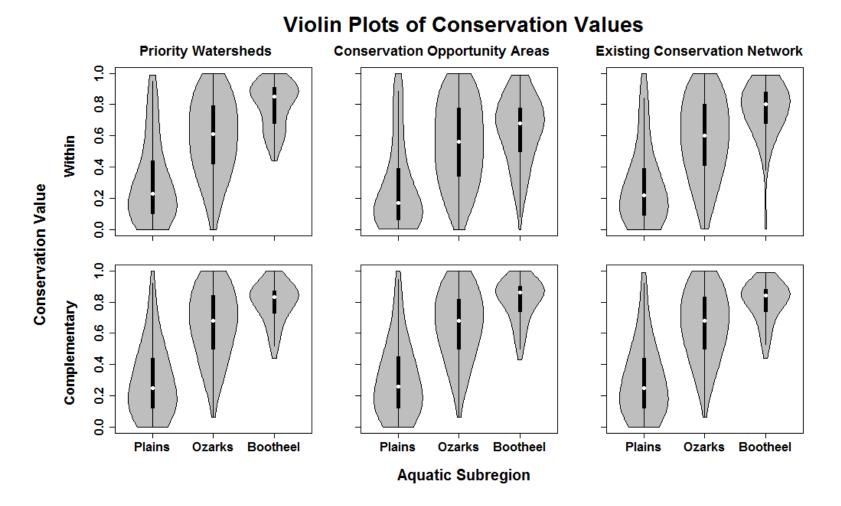


Figure 10: Violin plots of conservation value scores by subregion, separated by conservation network and within/complementary. The white point represents the median value, black bar represents the interquartile range, black line represents the overall range, and gray shaded area is a kernel density plot.

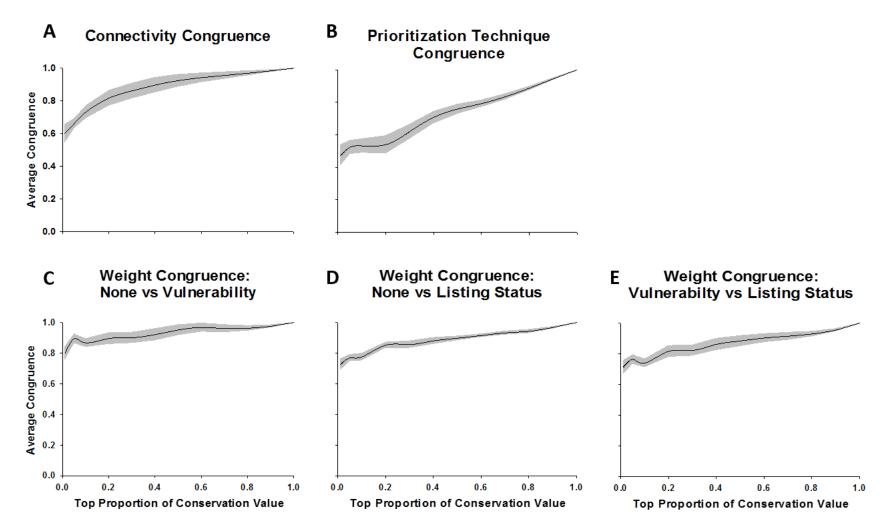


Figure 11: Plot of average percent congruence for the top proportion of conservation values with and without upstream connectivity requirements (A), each prioritization technique (B), and pairwise comparisons of each of the vulnerability approaches (C, D, E). Shaded areas represent the 95% confidence interval.

CONCLUSION

Our results may be used to support the management and conservation of stream fish in Missouri and can provide a framework for assessments in other regions. Our research suggested that the degree to which stream fish are vulnerable to climate change is likely to vary among species, and that different approaches to classifying vulnerability are likely to achieve different results (Chapter 1). We also found that high priority areas for stream fish conservation could be identified both within and complementary to established conservation networks (Chapter 2). We believe the ability to identify vulnerable fish species and areas which are important for their conservation is critical to the protection of stream fish species.

Our analysis of stream fish vulnerability revealed that although we have a relatively strong understanding of the role habitat degradation plays in species vulnerability, there is a great need for additional research to examine the effects of stream warming and alterations of flow regimes on stream fish. Our comparison of trait versus measured response-based approaches to assigning environmental tolerances yielded substantial discrepancies. Currently, the lack of data available for measuring species responses to climate related stressors limits the broad applicability and increases the uncertainty of a response based approach. The ability to extrapolate trait based sensitivities to all species, for which traits are known, makes the applicability of a trait association approach much more feasible. In addition, the mechanistic nature of a trait association approach provides logical support for assigning environmental

tolerances. However, additional factors not accounted for by traits may affect the environmental tolerances of some species. Based on currently available information for stream fish, we believe that the trait association index provides the most complete and reliable assessment of stream fish vulnerability. However, future research linking traits with environmental tolerances or measuring species responses to stressors is critical to improving the accuracy and decreasing the uncertainty of vulnerability assessments for stream fish.

The ability to assess the representation of stream fish within conservation networks is critical to developing effective conservation strategies. We found that by targeting areas within and complementary to established conservation networks, locations which provided comprehensive and redundant species representation could be identified. The comparison of results from our approach to selecting priority areas versus the use of surrogates, such as habitat factors or threat scores, revealed substantial increases in the predicted representation of species, particularly species which are rare or have restricted distributions, when our approach was used. The use of different prioritization techniques, species weighting, and inclusion of connectivity requirements had variable influences on the results of prioritization, suggesting careful consideration of inputs and settings is strongly advisable. Our comparison of species representation within the each of the conservation networks compared to an optimal network of the same size, generated using a blank slate approach, revealed that the proportion of representation lost for the species which lost the greatest proportion of its original distribution was the same or slightly higher when using core-area Zonation,

but lower when using the additive benefit function. The same comparison for the average proportion of distribution lost for all species revealed that for both core-area and additive benefit implementations species lost more representation when restricted to established networks than optimal networks of the same size. This suggests that predicted representations of species could be improved if conservation networks were developed from scratch although the relatively small differences suggest that these gains would be marginal and the benefits of working within established infrastructure likely outweigh the gains of starting from a blank slate. We believe that conservation value rankings of stream segments within conservation networks can be used to help prioritize the selection of sites for management actions. The conservation value rankings of stream segments complementary to established conservation networks can be used to aid in decisions related to land acquisition and the establishment of partnerships. Our framework for assessing stream segment scale conservation value while incorporating conservation networks which have already been established could be applied around the globe to assist in meeting the prioritization needs of managers and decision makers.

Our work also revealed a number of areas which would benefit greatly from additional research. One of the greatest needs is the development of stream temperature and flow models which represent both current and future conditions. Without this information the options for assessing the exposure of stream fish to climate change are extremely limited and riddled with uncertainty. Stream fish vulnerability assessments would also benefit greatly from research which quantifies the

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relationships between species or traits with stream temperature and flow regimes. Data on these topics for stream fish is currently very limited, making it difficult to draw conclusions as to the impact stressors will have on stream fish communities in the future. In order to more effectively plan for the conservation of stream fish species it would be beneficial to gain a greater understanding of how the landscape affects stream fish. It would be particularly useful to gain a better understanding of species-specific habitat requirements, specifically developing better measures of how upstream habitat degradation impacts a species, the area of suitable habitat which needs to be maintained for the persistence of a population (i.e. How big do protected areas need to be?) and how do factors such as connectivity dispersal ability, and metapopulation dynamics impact the viability of populations (i.e. Can a patchwork of sites protect a population and if so how close together do these sites need to be?).

APPENDICES

| Appendix 1: Species vulnerability scores for the response and trait based approaches to |
|---|
| classifying environmental tolerance |

| Species | Score (Response) | Score (Traits) |
|------------------------------|------------------|----------------|
| AMBLOPLITES ARIOMMUS | 3.12 | 4.65 |
| AMBLOPLITES CONSTELLATUS | 3.96 | 5.56 |
| AMBLOPLITES RUPESTRIS | 3.2 | 2.2 |
| AMEIURUS MELAS | 1.69 | 2.69 |
| AMEIURUS NATALIS | 0.6 | 1.6 |
| AMEIURUS NEBULOSUS | 2.87 | 4.86 |
| APHREDODERUS SAYANUS | 1.92 | 1.92 |
| APLODINOTUS GRUNNIENS | 0.94 | 2.48 |
| CAMPOSTOMA ANOMALUM PULLUM | 1.71 | 0.71 |
| CAMPOSTOMA OLIGOLEPIS | 1.01 | 2.01 |
| CARPIODES CARPIO | 1.05 | 2.49 |
| CARPIODES CYPRINUS | 1.18 | 2.83 |
| CARPIODES VELIFER | 4.61 | 4.61 |
| CATOSTOMUS COMMERSONII | 3.79 | 3.79 |
| CENTRARCHUS MACROPTERUS | | 4.86 |
| COTTUS BAIRDII | 3 | 5 |
| COTTUS CAROLINAE | 3.04 | 4.04 |
| COTTUS HYPSELURUS | | 6.22 |
| CYPRINELLA CAMURA | 2.47 | 2.47 |
| CYPRINELLA GALACTURA | 3.98 | 2.17 |
| CYPRINELLA LUTRENSIS | 0.86 | 0.86 |
| CYPRINELLA SPILOPTERA | 5.82 | 1.98 |
| CYPRINELLA VENUSTA | 1.67 | 3.42 |
| CYPRINELLA WHIPPLEI | 3.51 | 1.89 |
| DOROSOMA CEPEDIANUM | 0.67 | 2.67 |
| ERIMYSTAX HARRYI | | 5.63 |
| ERIMYSTAX X-PUNCTATUS | | 5.63 |
| ERIMYZON OBLONGUS | 2.85 | 5.63 |
| ESOX AMERICANUS VERMICULATUS | | 4.06 |
| ESOX NIGER | 3.94 | 3.94 |
| ETHEOSTOMA BLENNIOIDES | 3.87 | 2.87 |
| ETHEOSTOMA BURRI | | 7.87 |
| ETHEOSTOMA CAERULEUM | 2.99 | 3.99 |
| ETHEOSTOMA CRAGINI | | 7.82 |
| ETHEOSTOMA EUZONUM | | 8.11 |
| ETHEOSTOMA FLABELLARE | 2.97 | 3.97 |
| ETHEOSTOMA GRACILE | | 7.15 |
| ETHEOSTOMA JULIAE | | 8.48 |

| Species | Score (Response) | Score (Traits) |
|-------------------------|------------------|----------------|
| ETHEOSTOMA MICROPERCA | | 7.71 |
| ETHEOSTOMA NIANGUAE | | 8.75 |
| ETHEOSTOMA NIGRUM | 2.94 | 2.94 |
| ETHEOSTOMA PROELIARE | | 7.34 |
| ETHEOSTOMA PUNCTULATUM | 3.34 | 4.34 |
| ETHEOSTOMA SPECTABILE | 1.88 | 3.88 |
| ETHEOSTOMA STIGMAEUM | 7.82 | 5.84 |
| ETHEOSTOMA TETRAZONUM | | 6.87 |
| ETHEOSTOMA UNIPORUM | | 7.72 |
| ETHEOSTOMA ZONALE | 3.74 | 3.74 |
| FUNDULUS CATENATUS | 2.88 | 1.88 |
| FUNDULUS DISPAR | | 6.71 |
| FUNDULUS NOTATUS | 1.88 | 2.88 |
| FUNDULUS OLIVACEUS | 1.93 | 3.93 |
| FUNDULUS SCIADICUS | | 3.17 |
| GAMBUSIA AFFINIS | | 0.62 |
| HYBOGNATHUS ARGYRITIS | | 1.89 |
| HYBOGNATHUS HANKINSONI | 3.47 | 3.47 |
| HYBOGNATHUS PLACITUS | | 1.91 |
| HYBOPSIS AMBLOPS | 2.56 | 2.56 |
| HYPENTELIUM NIGRICANS | 2.02 | 4.02 |
| ICHTHYOMYZON FOSSOR | | 6.44 |
| ICHTHYOMYZON GAGEI | | 4.66 |
| ICTALURUS PUNCTATUS | 0.63 | 1.63 |
| ICTIOBUS BUBALUS | 1.15 | 2.94 |
| ICTIOBUS CYPRINELLUS | 1.51 | 3.32 |
| LAMPETRA AEPYPTERA | 2.07 | 5.91 |
| LEPISOSTEUS OCULATUS | | 3.66 |
| LEPISOSTEUS OSSEUS | 0.82 | 2.22 |
| LEPISOSTEUS PLATOSTOMUS | 0.96 | 2.46 |
| LEPOMIS CYANELLUS | 0.61 | 1.61 |
| LEPOMIS GULOSUS | 1.93 | 1.93 |
| LEPOMIS HUMILIS | 0.7 | 1.7 |
| LEPOMIS MACROCHIRUS | 0.62 | 1.62 |
| LEPOMIS MEGALOTIS | 1.68 | 2.68 |
| LEPOMIS MICROLOPHUS | 0.91 | 2.21 |
| LEPOMIS MINIATUS | | 3.29 |
| LUXILUS CARDINALIS | 3.45 | 4.9 |
| LUXILUS CHRYSOCEPHALUS | 1.89 | 1.89 |
| LUXILUS CORNUTUS | 2.83 | 2.83 |
| LUXILUS PILSBRYI | 3.54 | 4.92 |
| LUXILUS ZONATUS | | 2.05 |
| LYTHRURUS UMBRATILIS | 0.66 | 0.66 |

| Species | Score (Response) | Score (Traits) |
|--------------------------|------------------|----------------|
| MACRHYBOPSIS STORERIANA | 1.71 | 5.69 |
| MICROPTERUS DOLOMIEU | 2.99 | 1.99 |
| MICROPTERUS PUNCTULATUS | 0.86 | 2.16 |
| MICROPTERUS SALMOIDES | 0.62 | 1.62 |
| MINYTREMA MELANOPS | 2.56 | 4.15 |
| MORONE CHRYSOPS | 1.77 | 3.69 |
| MOXOSTOMA ANISURUM | 3.83 | 7.55 |
| MOXOSTOMA CARINATUM | 4.19 | 6.12 |
| MOXOSTOMA DUQUESNEI | 2.04 | 3.04 |
| MOXOSTOMA ERYTHRURUM | 0.96 | 2.96 |
| MOXOSTOMA MACROLEPIDOTUM | 0.82 | 5.26 |
| NOCOMIS ASPER | 3.8 | 6.94 |
| NOCOMIS BIGUTTATUS | 2 | 4 |
| NOTEMIGONUS CRYSOLEUCAS | 0.7 | 1.7 |
| NOTROPIS ATHERINOIDES | 2.79 | 2.79 |
| NOTROPIS BOOPS | 2.03 | 3.03 |
| NOTROPIS BUCCATUS | 2.34 | 4.1 |
| NOTROPIS BUCHANANI | | 3.9 |
| NOTROPIS DORSALIS | 2.05 | 2.05 |
| NOTROPIS GREENEI | | 3.2 |
| NOTROPIS HETEROLEPIS | | 2.33 |
| NOTROPIS NUBILUS | 2 | 3 |
| NOTROPIS OZARCANUS | | 5.83 |
| NOTROPIS STRAMINEUS | 1.99 | 0.99 |
| NOTROPIS TELESCOPUS | 2.5 | 3.61 |
| NOTROPIS TEXANUS | | 1.87 |
| NOTROPIS VOLUCELLUS | 3.1 | 1.46 |
| NOTURUS ALBATER | | 4.68 |
| NOTURUS EXILIS | 1.93 | 3.93 |
| NOTURUS FLAVATER | | 6.16 |
| NOTURUS FLAVUS | 4.61 | 6.33 |
| NOTURUS GYRINUS | 1.57 | 3.3 |
| NOTURUS MIURUS | 2.2 | 4.11 |
| NOTURUS NOCTURNUS | 1.4 | 7.34 |
| OPSOPOEODUS EMILIAE | | 1.34 |
| PERCINA CAPRODES | 2.79 | 2.79 |
| PERCINA COPELANDI | 4.43 | 6.4 |
| PERCINA CYMATOTAENIA | | 7.88 |
| PERCINA EVIDES | 5.98 | 7.92 |
| PERCINA MACULATA | 6.17 | 6.17 |
| PERCINA PHOXOCEPHALA | 4.76 | 6.39 |
| PERCINA SCIERA | 3.9 | 5.89 |
| PHENACOBIUS MIRABILIS | 1.84 | 1.84 |

| Species | Score (Response) | Score (Traits) |
|-------------------------|------------------|----------------|
| PHOXINUS ERYTHROGASTER | 1.95 | 1.95 |
| PIMEPHALES NOTATUS | 1.61 | 0.61 |
| PIMEPHALES PROMELAS | 1.64 | 0.64 |
| PIMEPHALES VIGILAX | 1.66 | 1.66 |
| POMOXIS ANNULARIS | 0.62 | 1.97 |
| POMOXIS NIGROMACULATUS | 1.17 | 3.04 |
| PYLODICTIS OLIVARIS | 0.96 | 2.6 |
| SANDER VITREUS | 4.21 | 8.19 |
| SEMOTILUS ATROMACULATUS | 0.65 | 2.65 |

| Species | Habitat | Thermal | Flow |
|------------------------------|---------|---------|------|
| AMBLOPLITES ARIOMMUS | 0 | 1 | 1 |
| AMBLOPLITES CONSTELLATUS | 0 | 1 | 1 |
| AMBLOPLITES RUPESTRIS | 0 | 0 | 1 |
| AMEIURUS MELAS | 1 | 0 | 1 |
| AMEIURUS NATALIS | 0 | 0 | 1 |
| AMEIURUS NEBULOSUS | 0 | 0 | 1 |
| APHREDODERUS SAYANUS | 0 | 0 | 0 |
| APLODINOTUS GRUNNIENS | 0 | 0 | 1 |
| CAMPOSTOMA ANOMALUM PULLUM | 0 | 0 | 0 |
| CAMPOSTOMA OLIGOLEPIS | 1 | 0 | 0 |
| CARPIODES CARPIO | 0 | 0 | 1 |
| CARPIODES CYPRINUS | 0 | 0 | 1 |
| CARPIODES VELIFER | 0 | 0 | 1 |
| CATOSTOMUS COMMERSONII | 1 | 1 | 1 |
| CENTRARCHUS MACROPTERUS | 0 | 0 | 1 |
| COTTUS BAIRDII | 1 | 1 | 1 |
| COTTUS CAROLINAE | 1 | 0 | 1 |
| COTTUS HYPSELURUS | 1 | 1 | 1 |
| CYPRINELLA CAMURA | 0 | 0 | 0 |
| CYPRINELLA GALACTURA | 0 | 0 | 0 |
| CYPRINELLA LUTRENSIS | 0 | 0 | 0 |
| CYPRINELLA SPILOPTERA | 0 | 0 | 0 |
| CYPRINELLA VENUSTA | 0 | 1 | 0 |
| CYPRINELLA WHIPPLEI | 0 | 0 | 0 |
| DOROSOMA CEPEDIANUM | 0 | 1 | 1 |
| ERIMYSTAX HARRYI | 1 | 1 | 0 |
| ERIMYSTAX X-PUNCTATUS | 1 | 1 | 0 |
| ERIMYZON OBLONGUS | 1 | 1 | 1 |
| ESOX AMERICANUS VERMICULATUS | 0 | 0 | 1 |
| ESOX NIGER | 0 | 0 | 1 |
| ETHEOSTOMA BLENNIOIDES | 1 | 0 | 0 |
| ETHEOSTOMA BURRI | 1 | 1 | 0 |
| ETHEOSTOMA CAERULEUM | 1 | 1 | 0 |
| ETHEOSTOMA CRAGINI | 1 | 1 | 0 |
| ETHEOSTOMA EUZONUM | 1 | 1 | 0 |
| ETHEOSTOMA FLABELLARE | 1 | 1 | 0 |
| ETHEOSTOMA GRACILE | 1 | 1 | 0 |
| ETHEOSTOMA JULIAE | 1 | 1 | 0 |
| ETHEOSTOMA MICROPERCA | 1 | 1 | 0 |

Appendix 2: Environmental tolerance classifications for the trait association index (1 = vulnerable, 0 = not vulnerable)

| Species | Habitat | Thermal | Flow |
|-------------------------|---------|---------|------|
| ETHEOSTOMA NIANGUAE | 1 | 1 | 0 |
| ETHEOSTOMA NIGRUM | 1 | 0 | 0 |
| ETHEOSTOMA PROELIARE | 1 | 1 | 0 |
| ETHEOSTOMA PUNCTULATUM | 1 | 1 | 0 |
| ETHEOSTOMA SPECTABILE | 1 | 1 | 0 |
| ETHEOSTOMA STIGMAEUM | 1 | 0 | 0 |
| ETHEOSTOMA TETRAZONUM | 1 | 1 | 0 |
| ETHEOSTOMA UNIPORUM | 1 | 1 | 0 |
| ETHEOSTOMA ZONALE | 1 | 0 | 0 |
| FUNDULUS CATENATUS | 0 | 0 | 0 |
| FUNDULUS DISPAR | 0 | 1 | 0 |
| FUNDULUS NOTATUS | 0 | 1 | 0 |
| FUNDULUS OLIVACEUS | 1 | 1 | 0 |
| FUNDULUS SCIADICUS | 0 | 0 | 0 |
| GAMBUSIA AFFINIS | 0 | 0 | 0 |
| HYBOGNATHUS ARGYRITIS | 0 | 0 | 0 |
| HYBOGNATHUS HANKINSONI | 0 | 1 | 0 |
| HYBOGNATHUS PLACITUS | 0 | 0 | 0 |
| HYBOPSIS AMBLOPS | 1 | 0 | 0 |
| HYPENTELIUM NIGRICANS | 1 | 1 | 1 |
| ICHTHYOMYZON FOSSOR | 1 | 1 | 0 |
| ICHTHYOMYZON GAGEI | 1 | 0 | 0 |
| ICTALURUS PUNCTATUS | 0 | 0 | 1 |
| ICTIOBUS BUBALUS | 0 | 0 | 1 |
| ICTIOBUS CYPRINELLUS | 0 | 0 | 1 |
| LAMPETRA AEPYPTERA | 1 | 1 | 0 |
| LEPISOSTEUS OCULATUS | 0 | 0 | 1 |
| LEPISOSTEUS OSSEUS | 0 | 0 | 1 |
| LEPISOSTEUS PLATOSTOMUS | 0 | 0 | 1 |
| LEPOMIS CYANELLUS | 0 | 0 | 1 |
| LEPOMIS GULOSUS | 0 | 0 | 1 |
| LEPOMIS HUMILIS | 1 | 0 | 0 |
| LEPOMIS MACROCHIRUS | 0 | 0 | 1 |
| LEPOMIS MEGALOTIS | 0 | 1 | 1 |
| LEPOMIS MICROLOPHUS | 0 | 0 | 1 |
| LEPOMIS MINIATUS | 0 | 0 | 1 |
| LUXILUS CARDINALIS | 1 | 1 | 0 |
| LUXILUS CHRYSOCEPHALUS | 1 | 0 | 0 |
| LUXILUS CORNUTUS | 1 | 0 | 0 |
| LUXILUS PILSBRYI | 1 | 1 | 0 |
| LUXILUS ZONATUS | 1 | 0 | 0 |
| LYTHRURUS UMBRATILIS | 0 | 0 | 0 |

| Species | Habitat | Thermal | Flow |
|--------------------------|---------|---------|------|
| MACRHYBOPSIS STORERIANA | 1 | 1 | 0 |
| MICROPTERUS DOLOMIEU | 0 | 0 | 1 |
| MICROPTERUS PUNCTULATUS | 0 | 0 | 1 |
| MICROPTERUS SALMOIDES | 0 | 0 | 1 |
| MINYTREMA MELANOPS | 1 | 0 | 1 |
| MORONE CHRYSOPS | 0 | 0 | 1 |
| MOXOSTOMA ANISURUM | 1 | 1 | 1 |
| MOXOSTOMA CARINATUM | 1 | 0 | 1 |
| MOXOSTOMA DUQUESNEI | 1 | 0 | 1 |
| MOXOSTOMA ERYTHRURUM | 1 | 0 | 1 |
| MOXOSTOMA MACROLEPIDOTUM | 1 | 1 | 1 |
| NOCOMIS ASPER | 1 | 1 | 1 |
| NOCOMIS BIGUTTATUS | 1 | 1 | 1 |
| NOTEMIGONUS CRYSOLEUCAS | 0 | 0 | 1 |
| NOTROPIS ATHERINOIDES | 0 | 1 | 0 |
| NOTROPIS BOOPS | 1 | 1 | 0 |
| NOTROPIS BUCCATUS | 1 | 0 | 0 |
| NOTROPIS BUCHANANI | 1 | 0 | 0 |
| NOTROPIS DORSALIS | 0 | 1 | 0 |
| NOTROPIS GREENEI | 1 | 0 | 0 |
| NOTROPIS HETEROLEPIS | 0 | 0 | 0 |
| NOTROPIS NUBILUS | 1 | 1 | 0 |
| NOTROPIS OZARCANUS | 1 | 1 | 0 |
| NOTROPIS STRAMINEUS | 0 | 0 | 0 |
| NOTROPIS TELESCOPUS | 1 | 1 | 0 |
| NOTROPIS TEXANUS | 0 | 0 | 0 |
| NOTROPIS VOLUCELLUS | 0 | 0 | 0 |
| NOTURUS ALBATER | 1 | 0 | 1 |
| NOTURUS EXILIS | 1 | 1 | 1 |
| NOTURUS FLAVATER | 1 | 0 | 1 |
| NOTURUS FLAVUS | 1 | 1 | 1 |
| NOTURUS GYRINUS | 1 | 0 | 0 |
| NOTURUS MIURUS | 1 | 0 | 0 |
| NOTURUS NOCTURNUS | 1 | 1 | 1 |
| OPSOPOEODUS EMILIAE | 0 | 0 | 0 |
| PERCINA CAPRODES | 1 | 0 | 0 |
| PERCINA COPELANDI | 1 | 0 | 0 |
| PERCINA CYMATOTAENIA | 1 | 1 | 0 |
| PERCINA EVIDES | 1 | 1 | 0 |
| PERCINA MACULATA | 1 | 1 | 0 |
| PERCINA PHOXOCEPHALA | 1 | 1 | 0 |
| PERCINA SCIERA | 1 | 0 | 0 |

| Species | Habitat | Thermal | Flow |
|-------------------------|---------|---------|------|
| PHENACOBIUS MIRABILIS | 1 | 0 | 0 |
| PHOXINUS ERYTHROGASTER | 0 | 1 | 0 |
| PIMEPHALES NOTATUS | 0 | 0 | 0 |
| PIMEPHALES PROMELAS | 0 | 0 | 0 |
| PIMEPHALES VIGILAX | 0 | 0 | 0 |
| POMOXIS ANNULARIS | 0 | 0 | 1 |
| POMOXIS NIGROMACULATUS | 0 | 0 | 1 |
| PYLODICTIS OLIVARIS | 0 | 0 | 1 |
| SANDER VITREUS | 0 | 1 | 1 |
| SEMOTILUS ATROMACULATUS | 1 | 1 | 0 |

| Species | Habitat | Thermal | Flow |
|------------------------------|---------|---------|------|
| AMBLOPLITES ARIOMMUS | 1 | 0 | 0 |
| AMBLOPLITES CONSTELLATUS | 1 | 0 | 0 |
| AMBLOPLITES RUPESTRIS | 1 | 0 | 1 |
| AMEIURUS MELAS | 0 | 0 | 1 |
| AMEIURUS NATALIS | 0 | 0 | 0 |
| AMEIURUS NEBULOSUS | 0 | 0 | 0 |
| APHREDODERUS SAYANUS | 0 | 0 | 0 |
| APLODINOTUS GRUNNIENS | 0 | 0 | 0 |
| CAMPOSTOMA ANOMALUM PULLUM | 0 | 1 | 0 |
| CAMPOSTOMA OLIGOLEPIS | 0 | 0 | 0 |
| CARPIODES CARPIO | 0 | 0 | 0 |
| CARPIODES CYPRINUS | 0 | 0 | 0 |
| CARPIODES VELIFER | 1 | 0 | 0 |
| CATOSTOMUS COMMERSONII | 1 | 1 | 1 |
| CENTRARCHUS MACROPTERUS | | | |
| COTTUS BAIRDII | 0 | 0 | 1 |
| COTTUS CAROLINAE | 1 | 0 | 0 |
| COTTUS HYPSELURUS | | | |
| CYPRINELLA CAMURA | 0 | 0 | 0 |
| CYPRINELLA GALACTURA | 1 | 0 | 0 |
| CYPRINELLA LUTRENSIS | 0 | 0 | 0 |
| CYPRINELLA SPILOPTERA | 1 | 0 | 1 |
| CYPRINELLA VENUSTA | 0 | 0 | 0 |
| CYPRINELLA WHIPPLEI | 1 | 0 | 0 |
| DOROSOMA CEPEDIANUM | 0 | 0 | 0 |
| ERIMYSTAX HARRYI | | | |
| ERIMYSTAX X-PUNCTATUS | | | |
| ERIMYZON OBLONGUS | 1 | 0 | 0 |
| ESOX AMERICANUS VERMICULATUS | | | |
| ESOX NIGER | 1 | 0 | 0 |
| ETHEOSTOMA BLENNIOIDES | 1 | 0 | 1 |
| ETHEOSTOMA BURRI | | | |
| ETHEOSTOMA CAERULEUM | 1 | 0 | 0 |
| ETHEOSTOMA CRAGINI | | | |
| ETHEOSTOMA EUZONUM | | | |
| ETHEOSTOMA FLABELLARE | 1 | 0 | 0 |
| ETHEOSTOMA GRACILE | | | |
| ETHEOSTOMA JULIAE | | | |
| ETHEOSTOMA MICROPERCA | | | |

Appendix 3: Environmental tolerance classifications for species response index (1 = vulnerable, 0 = not vulnerable)

| Species | Habitat | Thermal | Flow |
|-------------------------|---------|---------|------|
| ETHEOSTOMA NIANGUAE | | | |
| ETHEOSTOMA NIGRUM | 0 | 0 | 1 |
| ETHEOSTOMA PROELIARE | | | |
| ETHEOSTOMA PUNCTULATUM | 1 | 0 | 0 |
| ETHEOSTOMA SPECTABILE | 0 | 0 | 0 |
| ETHEOSTOMA STIGMAEUM | 1 | 0 | 1 |
| ETHEOSTOMA TETRAZONUM | | | |
| ETHEOSTOMA UNIPORUM | | | |
| ETHEOSTOMA ZONALE | 1 | 0 | 0 |
| FUNDULUS CATENATUS | 1 | 0 | 0 |
| FUNDULUS DISPAR | | | |
| FUNDULUS NOTATUS | 0 | 0 | 0 |
| FUNDULUS OLIVACEUS | 0 | 0 | 0 |
| FUNDULUS SCIADICUS | | | |
| GAMBUSIA AFFINIS | | | |
| HYBOGNATHUS ARGYRITIS | | | |
| HYBOGNATHUS HANKINSONI | 0 | 0 | 1 |
| HYBOGNATHUS PLACITUS | | | |
| HYBOPSIS AMBLOPS | 1 | 0 | 0 |
| HYPENTELIUM NIGRICANS | 1 | 0 | 0 |
| ICHTHYOMYZON FOSSOR | | | |
| ICHTHYOMYZON GAGEI | | | |
| ICTALURUS PUNCTATUS | 0 | 0 | 0 |
| ICTIOBUS BUBALUS | 0 | 0 | 0 |
| ICTIOBUS CYPRINELLUS | 0 | 0 | 0 |
| LAMPETRA AEPYPTERA | 0 | 0 | 0 |
| LEPISOSTEUS OCULATUS | | | |
| LEPISOSTEUS OSSEUS | 0 | 0 | 0 |
| LEPISOSTEUS PLATOSTOMUS | 0 | 0 | 0 |
| LEPOMIS CYANELLUS | 0 | 0 | 0 |
| LEPOMIS GULOSUS | 1 | 0 | 0 |
| LEPOMIS HUMILIS | 0 | 0 | 0 |
| LEPOMIS MACROCHIRUS | 0 | 0 | 0 |
| LEPOMIS MEGALOTIS | 1 | 0 | 0 |
| LEPOMIS MICROLOPHUS | 0 | 0 | 0 |
| LEPOMIS MINIATUS | | | |
| LUXILUS CARDINALIS | 1 | 0 | 0 |
| LUXILUS CHRYSOCEPHALUS | 1 | 0 | 0 |
| LUXILUS CORNUTUS | 0 | 0 | 1 |
| LUXILUS PILSBRYI | 1 | 0 | 0 |
| LUXILUS ZONATUS | | | |
| LYTHRURUS UMBRATILIS | 0 | 0 | 0 |

| Species | Habitat | Thermal | Flow |
|--------------------------|---------|---------|------|
| MACRHYBOPSIS STORERIANA | 0 | 0 | 0 |
| MICROPTERUS DOLOMIEU | 1 | 1 | 0 |
| MICROPTERUS PUNCTULATUS | 0 | 0 | 0 |
| MICROPTERUS SALMOIDES | 0 | 0 | 0 |
| MINYTREMA MELANOPS | 1 | 0 | 0 |
| MORONE CHRYSOPS | 0 | 0 | 0 |
| MOXOSTOMA ANISURUM | 0 | 0 | 1 |
| MOXOSTOMA CARINATUM | 1 | 0 | 0 |
| MOXOSTOMA DUQUESNEI | 1 | 0 | 0 |
| MOXOSTOMA ERYTHRURUM | 0 | 0 | 0 |
| MOXOSTOMA MACROLEPIDOTUM | 0 | 0 | 0 |
| NOCOMIS ASPER | 1 | 0 | 0 |
| NOCOMIS BIGUTTATUS | 1 | 0 | 0 |
| NOTEMIGONUS CRYSOLEUCAS | 0 | 0 | 0 |
| NOTROPIS ATHERINOIDES | 0 | 0 | 1 |
| NOTROPIS BOOPS | 1 | 0 | 0 |
| NOTROPIS BUCCATUS | 0 | 0 | 0 |
| NOTROPIS BUCHANANI | | | |
| NOTROPIS DORSALIS | 0 | 0 | 1 |
| NOTROPIS GREENEI | | | |
| NOTROPIS HETEROLEPIS | | | |
| NOTROPIS NUBILUS | 1 | 0 | 0 |
| NOTROPIS OZARCANUS | | | |
| NOTROPIS STRAMINEUS | 1 | 0 | 0 |
| NOTROPIS TELESCOPUS | 1 | 0 | 0 |
| NOTROPIS TEXANUS | | | |
| NOTROPIS VOLUCELLUS | 0 | 0 | 1 |
| NOTURUS ALBATER | | | |
| NOTURUS EXILIS | 1 | 0 | 0 |
| NOTURUS FLAVATER | | | |
| NOTURUS FLAVUS | 1 | 0 | 1 |
| NOTURUS GYRINUS | 0 | 0 | 0 |
| NOTURUS MIURUS | 0 | 0 | 0 |
| NOTURUS NOCTURNUS | 0 | 0 | 0 |
| OPSOPOEODUS EMILIAE | | | |
| PERCINA CAPRODES | 1 | 0 | 0 |
| PERCINA COPELANDI | 0 | 0 | 0 |
| PERCINA CYMATOTAENIA | | | |
| PERCINA EVIDES | 1 | 0 | 0 |
| PERCINA MACULATA | 1 | 0 | 1 |
| PERCINA PHOXOCEPHALA | 1 | 0 | 0 |
| PERCINA SCIERA | 0 | 0 | 0 |

| Species | Habitat | Thermal | Flow |
|-------------------------|---------|---------|------|
| PHENACOBIUS MIRABILIS | 1 | 0 | 0 |
| PHOXINUS ERYTHROGASTER | 1 | 0 | 0 |
| PIMEPHALES NOTATUS | 0 | 0 | 1 |
| PIMEPHALES PROMELAS | 0 | 0 | 1 |
| PIMEPHALES VIGILAX | 0 | 0 | 0 |
| POMOXIS ANNULARIS | 0 | 0 | 0 |
| POMOXIS NIGROMACULATUS | 0 | 0 | 0 |
| PYLODICTIS OLIVARIS | 0 | 0 | 0 |
| SANDER VITREUS | 0 | 0 | 0 |
| SEMOTILUS ATROMACULATUS | 0 | 0 | 0 |

| Species | Rarity | Dispersal | Range | Resilience |
|------------------------------|--------|-----------|-------|------------|
| AMBLOPLITES ARIOMMUS | 1.53 | 0 | 0.765 | 0.42 |
| AMBLOPLITES CONSTELLATUS | 1.6 | 0 | 0.942 | 0.85 |
| AMBLOPLITES RUPESTRIS | 1 | 0 | 0.589 | 0.61 |
| AMEIURUS MELAS | 1 | 0 | 0.177 | 0.51 |
| AMEIURUS NATALIS | 1 | 0 | 0 | 0.6 |
| AMEIURUS NEBULOSUS | 1.99 | 0 | 0.942 | 1 |
| APHREDODERUS SAYANUS | 1.82 | 0 | 0.765 | 0.53 |
| APLODINOTUS GRUNNIENS | 1.54 | 0 | 0.236 | 0.58 |
| CAMPOSTOMA ANOMALUM PULLUM | 1 | 0 | 0.118 | 0.59 |
| CAMPOSTOMA OLIGOLEPIS | 1 | 0 | 0.412 | 0.6 |
| CARPIODES CARPIO | 1.44 | 0 | 0.471 | 0.37 |
| CARPIODES CYPRINUS | 1.65 | 0 | 0.412 | 0.5 |
| CARPIODES VELIFER | 1.98 | 0 | 0.883 | 0.88 |
| CATOSTOMUS COMMERSONII | 1 | 0 | 0.177 | 0.61 |
| CENTRARCHUS MACROPTERUS | 1.99 | 0 | 0.942 | 1 |
| COTTUS BAIRDII | 1 | 1 | 0.471 | 0.53 |
| COTTUS CAROLINAE | 1 | 1 | 0.412 | 0.63 |
| COTTUS HYPSELURUS | 1.22 | 1 | 0.648 | 0.55 |
| CYPRINELLA CAMURA | 1.99 | 0 | 0.942 | 0.6 |
| CYPRINELLA GALACTURA | 1.81 | 0 | 0.824 | 0.68 |
| CYPRINELLA LUTRENSIS | 1 | 0 | 0.295 | 0.56 |
| CYPRINELLA SPILOPTERA | 1.92 | 0 | 0.706 | 0.62 |
| CYPRINELLA VENUSTA | 1.75 | 0 | 0.648 | 0.54 |
| CYPRINELLA WHIPPLEI | 1.62 | 0 | 0.765 | 0.65 |
| DOROSOMA CEPEDIANUM | 1 | 0 | 0.059 | 0.61 |
| ERIMYSTAX HARRYI | 1.87 | 0 | 0.824 | 0.35 |
| ERIMYSTAX X-PUNCTATUS | 1.9 | 0 | 0.589 | 0.71 |
| ERIMYZON OBLONGUS | 1.39 | 0 | 0.648 | 0.56 |
| ESOX AMERICANUS VERMICULATUS | 1.85 | 0 | 0.765 | 0.79 |
| ESOX NIGER | 1.86 | 0 | 0.765 | 0.66 |
| ETHEOSTOMA BLENNIOIDES | 1 | 1 | 0.236 | 0.63 |
| ETHEOSTOMA BURRI | 1.76 | 1 | 0.942 | 0.93 |
| ETHEOSTOMA CAERULEUM | 1 | 1 | 0.353 | 0.64 |
| ETHEOSTOMA CRAGINI | 1.82 | 1 | 0.942 | 0.65 |
| ETHEOSTOMA EUZONUM | 1.99 | 1 | 0.942 | 0.27 |
| ETHEOSTOMA FLABELLARE | 1 | 1 | 0.295 | 0.67 |
| ETHEOSTOMA GRACILE | 1.94 | 1 | 0.589 | 0.19 |
| ETHEOSTOMA JULIAE | 1.93 | 1 | 0.942 | 0.87 |
| ETHEOSTOMA MICROPERCA | 1.97 | 1 | 0.824 | 0.18 |
| ETHEOSTOMA NIANGUAE | 1.99 | 1 | 0.942 | 0.91 |

Appendix 4: Species scores for rarity, dispersal, range and freshwater resilience

| Species | Rarity | Dispersal | Range | Resilience |
|-------------------------|--------|-----------|-------|------------|
| ETHEOSTOMA NIGRUM | 1 | 1 | 0.353 | 0.59 |
| ETHEOSTOMA PROELIARE | 1.95 | 1 | 0.765 | 0 |
| ETHEOSTOMA PUNCTULATUM | 1 | 1 | 0.648 | 0.69 |
| ETHEOSTOMA SPECTABILE | 1 | 1 | 0.236 | 0.64 |
| ETHEOSTOMA STIGMAEUM | 1.98 | 1 | 0.706 | 0.48 |
| ETHEOSTOMA TETRAZONUM | 1.68 | 1 | 0.706 | 0.64 |
| ETHEOSTOMA UNIPORUM | 1.95 | 1 | 0.942 | 0.03 |
| ETHEOSTOMA ZONALE | 1.26 | 1 | 0.471 | 0.63 |
| FUNDULUS CATENATUS | 1 | 1 | 0.236 | 0.64 |
| FUNDULUS DISPAR | 1.98 | 1 | 0.883 | 1 |
| FUNDULUS NOTATUS | 1 | 1 | 0.177 | 0.7 |
| FUNDULUS OLIVACEUS | 1 | 1 | 0.236 | 0.69 |
| FUNDULUS SCIADICUS | 1.72 | 1 | 0.648 | 0.34 |
| GAMBUSIA AFFINIS | 1 | 0 | 0 | 0.62 |
| HYBOGNATHUS ARGYRITIS | 1.97 | 0 | 0.883 | 0.15 |
| HYBOGNATHUS HANKINSONI | 1.78 | 0 | 0.883 | 0.12 |
| HYBOGNATHUS PLACITUS | 1.98 | 0 | 0.824 | 0.28 |
| HYBOPSIS AMBLOPS | 1.09 | 0 | 0.648 | 0.76 |
| HYPENTELIUM NIGRICANS | 1 | 0 | 0.353 | 0.67 |
| ICHTHYOMYZON FOSSOR | 1.99 | 0 | 0.883 | 0.7 |
| ICHTHYOMYZON GAGEI | 1.97 | 0 | 0.942 | 0.83 |
| ICTALURUS PUNCTATUS | 1 | 0 | 0.118 | 0.51 |
| ICTIOBUS BUBALUS | 1.79 | 0 | 0.353 | 0.52 |
| ICTIOBUS CYPRINELLUS | 1.81 | 0 | 0.412 | 0.76 |
| LAMPETRA AEPYPTERA | 1.92 | 0 | 0.883 | 0.37 |
| LEPISOSTEUS OCULATUS | 1.79 | 0 | 0.765 | 0.5 |
| LEPISOSTEUS OSSEUS | 1.4 | 0 | 0.177 | 0.57 |
| LEPISOSTEUS PLATOSTOMUS | 1.5 | 0 | 0.295 | 0.52 |
| LEPOMIS CYANELLUS | 1 | 0 | 0 | 0.61 |
| LEPOMIS GULOSUS | 1.19 | 0 | 0.177 | 0.53 |
| LEPOMIS HUMILIS | 1 | 0 | 0.118 | 0.58 |
| LEPOMIS MACROCHIRUS | 1 | 0 | 0 | 0.62 |
| LEPOMIS MEGALOTIS | 1 | 0 | 0 | 0.68 |
| LEPOMIS MICROLOPHUS | 1.3 | 0 | 0.177 | 0.68 |
| LEPOMIS MINIATUS | 1.63 | 0 | 0.706 | 0.51 |
| LUXILUS CARDINALIS | 1.45 | 0 | 0.883 | 0.72 |
| LUXILUS CHRYSOCEPHALUS | 1 | 0 | 0.295 | 0.59 |
| LUXILUS CORNUTUS | 1.27 | 0 | 0.706 | 0.66 |
| LUXILUS PILSBRYI | 1.38 | 0 | 0.942 | 0.86 |
| LUXILUS ZONATUS | 1 | 0 | 0.471 | 0.58 |
| LYTHRURUS UMBRATILIS | 1 | 0 | 0.177 | 0.48 |
| MACRHYBOPSIS STORERIANA | 1.99 | 0 | 0.706 | 0.31 |

| Species | Rarity | Dispersal | Range | Resilience |
|--------------------------|--------|-----------|-------|------------|
| MICROPTERUS DOLOMIEU | 1 | 0 | 0.353 | 0.64 |
| MICROPTERUS PUNCTULATUS | 1.3 | 0 | 0.118 | 0.71 |
| MICROPTERUS SALMOIDES | 1 | 0 | 0 | 0.62 |
| MINYTREMA MELANOPS | 1.59 | 0 | 0.295 | 0.5 |
| MORONE CHRYSOPS | 1.92 | 0 | 0.53 | 0.75 |
| MOXOSTOMA ANISURUM | 1.86 | 0 | 0.706 | 0.66 |
| MOXOSTOMA CARINATUM | 1.93 | 0 | 0.824 | 0.67 |
| MOXOSTOMA DUQUESNEI | 1 | 0 | 0.353 | 0.69 |
| MOXOSTOMA ERYTHRURUM | 1 | 0 | 0.236 | 0.72 |
| MOXOSTOMA MACROLEPIDOTUM | 1.48 | 0 | 0.118 | 0.65 |
| NOCOMIS ASPER | 1.57 | 0 | 0.942 | 0.75 |
| NOCOMIS BIGUTTATUS | 1 | 0 | 0.412 | 0.59 |
| NOTEMIGONUS CRYSOLEUCAS | 1 | 0 | 0.059 | 0.64 |
| NOTROPIS ATHERINOIDES | 1.65 | 0 | 0.295 | 0.65 |
| NOTROPIS BOOPS | 1 | 0 | 0.412 | 0.62 |
| NOTROPIS BUCCATUS | 1.76 | 0 | 0.824 | 0.89 |
| NOTROPIS BUCHANANI | 1.96 | 0 | 0.706 | 0.56 |
| NOTROPIS DORSALIS | 1 | 0 | 0.589 | 0.46 |
| NOTROPIS GREENEI | 1.59 | 0 | 0.648 | 0.58 |
| NOTROPIS HETEROLEPIS | 1.95 | 0 | 0.765 | 0.84 |
| NOTROPIS NUBILUS | 1 | 0 | 0.353 | 0.65 |
| NOTROPIS OZARCANUS | 1.93 | 0 | 0.824 | 0.38 |
| NOTROPIS STRAMINEUS | 1 | 0 | 0.412 | 0.58 |
| NOTROPIS TELESCOPUS | 1.11 | 0 | 0.765 | 0.54 |
| NOTROPIS TEXANUS | 1.86 | 0 | 0.824 | 0.34 |
| NOTROPIS VOLUCELLUS | 1.64 | 0 | 0.471 | 0.69 |
| NOTURUS ALBATER | 1.43 | 0 | 0.824 | 0.64 |
| NOTURUS EXILIS | 1 | 0 | 0.236 | 0.69 |
| NOTURUS FLAVATER | 1.98 | 0 | 0.883 | 0.45 |
| NOTURUS FLAVUS | 1.72 | 0 | 0.353 | 0.56 |
| NOTURUS GYRINUS | 1.73 | 0 | 0.589 | 0.55 |
| NOTURUS MIURUS | 1.91 | 0 | 0.824 | 0.63 |
| NOTURUS NOCTURNUS | 1.98 | 0 | 0.589 | 0.23 |
| OPSOPOEODUS EMILIAE | 1.89 | 0 | 0.706 | 0.01 |
| PERCINA CAPRODES | 1 | 1 | 0.118 | 0.67 |
| PERCINA COPELANDI | 1.97 | 1 | 0.942 | 0.6 |
| PERCINA CYMATOTAENIA | 1.98 | 1 | 0.883 | 0.19 |
| PERCINA EVIDES | 1.94 | 1 | 0.706 | 0.73 |
| PERCINA MACULATA | 1.59 | 1 | 0.53 | 0.56 |
| PERCINA PHOXOCEPHALA | 1.63 | 1 | 0.53 | 0.64 |
| PERCINA SCIERA | 1.99 | 1 | 0.765 | 0.39 |
| PHENACOBIUS MIRABILIS | 1 | 0 | 0.353 | 0.49 |

| Species | Rarity | Dispersal | Range | Resilience |
|-------------------------|--------|-----------|-------|------------|
| PHOXINUS ERYTHROGASTER | 1 | 0 | 0.353 | 0.6 |
| PIMEPHALES NOTATUS | 1 | 0 | 0 | 0.61 |
| PIMEPHALES PROMELAS | 1 | 0 | 0.177 | 0.46 |
| PIMEPHALES VIGILAX | 1.72 | 0 | 0.589 | 0.65 |
| POMOXIS ANNULARIS | 1.35 | 0 | 0.059 | 0.54 |
| POMOXIS NIGROMACULATUS | 1.87 | 0 | 0.236 | 0.73 |
| PYLODICTIS OLIVARIS | 1.64 | 0 | 0.295 | 0.48 |
| SANDER VITREUS | 1.99 | 1 | 0.765 | 0.7 |
| SEMOTILUS ATROMACULATUS | 1 | 0 | 0.059 | 0.59 |

| Variable | Units |
|---|-----------------------------------|
| Ecological Drainage Unit | Categorical |
| Aquatic Ecological System Type | Categorical |
| Strahler Stream Order | Continuous Categorical |
| Downstream Size | Continuous Categorical |
| Reach Gradient | m/km |
| Segment Gradient | m/km |
| Local Geological Classification | Categorical |
| Urban Landcover- Local | % of Local Watershed Area |
| Agricultural Landcover- Local | % of Local Watershed Area |
| Grassland Landcover- Local | % of Local Watershed Area |
| Forest Landcover- Local | % of Local Watershed Area |
| Wetland Landcover- Local | % of Local Watershed Area |
| Urban Landcover- Upstream Watershed | % of Upstream Watershed Area |
| Agricultural Landcover- Upstream Watershed | % of Upstream Watershed Area |
| Grassland Landcover- Upstream Watershed | % of Upstream Watershed Area |
| Forest Landcover- Upstream Watershed | % of Upstream Watershed Area |
| Wetland Landcover- Upstream Watershed | % of Upstream Watershed Area |
| Number of Springs- Local | Count |
| Number of Springs- Watershed | Count |
| Number of Dams- Local | Count |
| Number of Dams- Upstream Watershed | Count |
| Number of Headwater Impoundments- Local | Count |
| Number of Headwater Impoundments- Upstream | Count |
| Watershed | |
| Number of Road/Stream Crossings per KM ² - Local | Ratio |
| Number of Road/Stream Crossings per KM ² - | Ratio |
| Upstream Watershed | |
| Local Habitat Threat Index Score | Value |
| Upstream Habitat Threat Index Score | Value |
| Percent of the Local Riparian Intact | % of Local Riparian Not Classifie |
| | as Agriculture or Urban |

Appendix 5: Environmental variables and associated units which were used for the creation of species distribution models

Appendix 6: A list of species codes, scientific names, and common names for all study species, as well as whether a model was constructed for each species for both the Plains and Ozarks subregions. Blank indicates no model constructed, "Yes" indicates a satisfactory model was created, "Yes (Failed)" indicates a model was created but did not meet the model evaluation criteria.

| Species Code | Scientific Name | Plains Model | Ozark Model |
|----------------|--------------------------|--------------|-------------|
| A_ARIOMMUS | AMBLOPLITES ARIOMMUS | | Yes |
| A_CONSTELLATUS | AMBLOPLITES CONSTELLATUS | | Yes |
| A_GRUNNIENS | APLODINOTUS GRUNNIENS | Yes | |
| A_MELAS | AMEIURUS MELAS | Yes (Failed) | Yes |
| A_NATALIS | AMEIURUS NATALIS | Yes | Yes |
| A_NEBULOSUS | AMEIURUS NEBULOSUS | | |
| A_RUPESTRIS | AMBLOPLITES RUPESTRIS | | Yes |
| A_SAYANUS | APHREDODERUS SAYANUS | | |
| C_BAIRDII | COTTUS BAIRDII | | Yes |
| C_CAMURA | CYPRINELLA CAMURA | | |
| C_CAROLINAE | COTTUS CAROLINAE | | Yes |
| C_CARPIO | CARPIODES CARPIO | Yes | |
| C_COMMERSONII | CATOSTOMUS COMMERSONII | Yes | Yes |
| C_CYPRINUS | CARPIODES CYPRINUS | Yes | |
| C_GALACTURA | CYPRINELLA GALACTURA | | Yes |
| C_HYPSELURUS | COTTUS HYPSELURUS | | Yes |
| C_LUTRENSIS | CYPRINELLA LUTRENSIS | Yes | Yes |
| C_MACROPTERUS | CENTRARCHUS MACROPTERUS | | |
| C_OLIGOLEPIS | CAMPOSTOMA OLIGOLEPIS | | Yes |
| | CAMPOSTOMA ANOMALUM | | |
| C_PULLUM | PULLUM | Yes | Yes |
| C_SPILOPTERA | CYPRINELLA SPILOPTERA | | |
| C_VELIFER | CARPIODES VELIFER | | |
| C_VENUSTA | CYPRINELLA VENUSTA | | |
| C_WHIPPLEI | CYPRINELLA WHIPPLEI | | Yes |
| D_CEPEDIANUM | DOROSOMA CEPEDIANUM | Yes | Yes |
| E_BLENNIOIDES | ETHEOSTOMA BLENNIOIDES | | Yes |
| E_BURRI | ETHEOSTOMA BURRI | | |
| E_CAERULEUM | ETHEOSTOMA CAERULEUM | | Yes |
| E_CRAGINI | ETHEOSTOMA CRAGINI | | |
| E_EUZONUM | ETHEOSTOMA EUZONUM | | |
| E_FLABELLARE | ETHEOSTOMA FLABELLARE | Yes | Yes |
| E_GRACILE | ETHEOSTOMA GRACILE | | |
| E_HARRYI | ERIMYSTAX HARRYI | | |
| E_JULIAE | ETHEOSTOMA JULIAE | | |
| E_MICROPERCA | ETHEOSTOMA MICROPERCA | | |

| Species Code | Scientific Name | Plains Model | Ozark Model |
|-----------------|------------------------|---------------------|-------------|
| E_NIANGUAE | ETHEOSTOMA NIANGUAE | | |
| E_NIGER | ESOX NIGER | | |
| E_NIGRUM | ETHEOSTOMA NIGRUM | Yes | Yes |
| E_OBLONGUS | ERIMYZON OBLONGUS | | Yes |
| E_PROELIARE | ETHEOSTOMA PROELIARE | | |
| E_PUNCTULATUM | ETHEOSTOMA PUNCTULATUM | | Yes |
| E_SPECTABILE | ETHEOSTOMA SPECTABILE | Yes | Yes |
| e_stigmaeum | ETHEOSTOMA STIGMAEUM | | |
| E_TETRAZONUM | ETHEOSTOMA TETRAZONUM | | Yes |
| E_UNIPORUM | ETHEOSTOMA UNIPORUM | | |
| | ESOX AMERICANUS | | |
| E_VERMICULATUS | VERMICULATUS | | |
| E_X-PUNCTATUS | ERIMYSTAX X-PUNCTATUS | | |
| E_ZONALE | ETHEOSTOMA ZONALE | | Yes |
| F_CATENATUS | FUNDULUS CATENATUS | | Yes |
| F_DISPAR | FUNDULUS DISPAR | | |
| F_NOTATUS | FUNDULUS NOTATUS | Yes | Yes |
| F_OLIVACEUS | FUNDULUS OLIVACEUS | | Yes |
| F_SCIADICUS | FUNDULUS SCIADICUS | | Yes |
| G_AFFINIS | GAMBUSIA AFFINIS | Yes | Yes |
| H_AMBLOPS | HYBOPSIS AMBLOPS | | Yes |
| H_ARGYRITIS | HYBOGNATHUS ARGYRITIS | | |
| H_HANKINSONI | HYBOGNATHUS HANKINSONI | | |
| H_NIGRICANS | HYPENTELIUM NIGRICANS | | Yes |
| H_PLACITUS | HYBOGNATHUS PLACITUS | | |
| I_BUBALUS | ICTIOBUS BUBALUS | | |
| I_CYPRINELLUS | ICTIOBUS CYPRINELLUS | | |
| I_FOSSOR | ICHTHYOMYZON FOSSOR | | |
| I_GAGEI | ICHTHYOMYZON GAGEI | | |
| I_NIGER | ICTIOBUS NIGER | | |
| I_PUNCTATUS | ICTALURUS PUNCTATUS | Yes | Yes |
| L_AEPYPTERA | LAMPETRA AEPYPTERA | | |
| L_CARDINALIS | LUXILUS CARDINALIS | | Yes |
| L_CHRYSOCEPHALU | | | |
| S | LUXILUS CHRYSOCEPHALUS | Yes | Yes |
| L_CORNUTUS | LUXILUS CORNUTUS | Yes | Yes |
| L_CYANELLUS | LEPOMIS CYANELLUS | Yes (Failed) | Yes |
| L_GULOSUS | LEPOMIS GULOSUS | | Yes |
| L_HUMILIS | LEPOMIS HUMILIS | Yes | Yes |
| L_MACROCHIRUS | LEPOMIS MACROCHIRUS | Yes | Yes |
| L_MEGALOTIS | LEPOMIS MEGALOTIS | Yes | Yes |
| L_MICROLOPHUS | LEPOMIS MICROLOPHUS | | Yes |

| Species Code | Scientific Name | Plains Model | Ozark Model |
|-------------------|-------------------------|---------------------|-------------|
| L_MINIATUS | LEPOMIS MINIATUS | | |
| L_OCULATUS | LEPISOSTEUS OCULATUS | | |
| L_OSSEUS | LEPISOSTEUS OSSEUS | | Yes |
| L_PILSBRYI | LUXILUS PILSBRYI | | Yes |
| L_PLATOSTOMUS | LEPISOSTEUS PLATOSTOMUS | Yes | |
| L_UMBRATILIS | LYTHRURUS UMBRATILIS | Yes | Yes |
| L_ZONATUS | LUXILUS ZONATUS | | Yes |
| M_ANISURUM | MOXOSTOMA ANISURUM | | |
| M_CARINATUM | MOXOSTOMA CARINATUM | | |
| M_CHRYSOPS | MORONE CHRYSOPS | | |
| M_DOLOMIEU | MICROPTERUS DOLOMIEU | | Yes |
| M_DUQUESNEI | MOXOSTOMA DUQUESNEI | | Yes |
| M ERYTHRURUM | MOXOSTOMA ERYTHRURUM | Yes | Yes |
| M_MACROLEPIDOT | ΜΟΧΟΣΤΟΜΑ | | |
| UM | MACROLEPIDOTUM | | Yes |
| M_MELANOPS | MINYTREMA MELANOPS | | Yes |
| M_PUNCTULATUS | MICROPTERUS PUNCTULATUS | | Yes |
| M_SALMOIDES | MICROPTERUS SALMOIDES | Yes | Yes |
| M_STORERIANA | MACRHYBOPSIS STORERIANA | | |
| N_ALBATER | NOTURUS ALBATER | | Yes |
| N_ASPER | NOCOMIS ASPER | | Yes |
| N_ATHERINOIDES | NOTROPIS ATHERINOIDES | | |
| N_BIGUTTATUS | NOCOMIS BIGUTTATUS | | Yes |
| N_BOOPS | NOTROPIS BOOPS | Yes | Yes |
| N_BUCCATUS | NOTROPIS BUCCATUS | | |
| N_BUCHANANI | NOTROPIS BUCHANANI | | |
| N_CRYSOLEUCAS | NOTEMIGONUS CRYSOLEUCAS | Yes | Yes |
| N_DORSALIS | NOTROPIS DORSALIS | Yes | |
| N_EXILIS | NOTURUS EXILIS | Yes | Yes |
| N_FLAVATER | NOTURUS FLAVATER | | |
| N_FLAVUS | NOTURUS FLAVUS | Yes | |
| N_GREENEI | NOTROPIS GREENEI | | Yes |
| N GYRINUS | NOTURUS GYRINUS | | |
| N HETEROLEPIS | NOTROPIS HETEROLEPIS | | |
| N MIURUS | NOTURUS MIURUS | | |
| N_NOCTURNUS | NOTURUS NOCTURNUS | | |
| N NUBILUS | NOTROPIS NUBILUS | | Yes |
| N_OZARCANUS | NOTROPIS OZARCANUS | | |
| | NOTROPIS STRAMINEUS | Yes | Yes |
| _ N_TELESCOPUS | NOTROPIS TELESCOPUS | | Yes |
| _ N_TEXANUS | NOTROPIS TEXANUS | | |
| N VOLUCELLUS | NOTROPIS VOLUCELLUS | | |

| Species Code | Scientific Name | Plains Model | Ozark Model |
|-----------------|-------------------------|--------------|-------------|
| O_EMILIAE | OPSOPOEODUS EMILIAE | | |
| P_ANNULARIS | POMOXIS ANNULARIS | Yes | |
| P_CAPRODES | PERCINA CAPRODES | Yes | Yes |
| P_COPELANDI | PERCINA COPELANDI | | |
| P_CYMATOTAENIA | PERCINA CYMATOTAENIA | | |
| P_ERYTHROGASTER | PHOXINUS ERYTHROGASTER | | Yes |
| P_EVIDES | PERCINA EVIDES | | |
| P_FLAVESCENS | PERCA FLAVESCENS | | |
| P_MACULATA | PERCINA MACULATA | Yes | |
| P_MIRABILIS | PHENACOBIUS MIRABILIS | Yes | Yes |
| P_NIGROMACULAT | | | |
| US | POMOXIS NIGROMACULATUS | | |
| P_NOTATUS | PIMEPHALES NOTATUS | Yes | Yes |
| P_OLIVARIS | PYLODICTIS OLIVARIS | Yes | |
| P_PHOXOCEPHALA | PERCINA PHOXOCEPHALA | Yes | |
| P_PROMELAS | PIMEPHALES PROMELAS | Yes | Yes |
| P_SCIERA | PERCINA SCIERA | | |
| P_VIGILAX | PIMEPHALES VIGILAX | | |
| S_ATROMACULATU | | | |
| S | SEMOTILUS ATROMACULATUS | Yes | Yes |
| S_VITREUS | SANDER VITREUS | | |

Appendix 7: A list of component models which went into the ensemble for each species. Component models assigned a score of 0 did not meet minimum requirements for at least one evaluation statistic and therefore were not included in the ensemble, while component models assigned a score of 1 met minimum requirements for all three model evaluation statistics and were included in the final ensemble.

| Species | MARS | GAM | BRT | RndmFrst | Sum |
|----------------------------|------|-----|-----|----------|-----|
| AMEIURUS MELAS | 0 | 0 | 0 | 0 | 0 |
| AMEIURUS NATALIS | 0 | 0 | 1 | 1 | 2 |
| APLODINOTUS GRUNNIENS | 0 | 0 | 0 | 1 | 1 |
| CAMPOSTOMA ANOMALUM PULLUM | 0 | 1 | 1 | 1 | 3 |
| CARPIODES CARPIO | 1 | 1 | 1 | 1 | 4 |
| CARPIODES CYPRINUS | 1 | 1 | 0 | 1 | 3 |
| CATOSTOMUS COMMERSONII | 0 | 1 | 0 | 1 | 2 |
| CYPRINELLA LUTRENSIS | 1 | 1 | 1 | 1 | 4 |
| DOROSOMA CEPEDIANUM | 0 | 1 | 0 | 1 | 2 |
| ETHEOSTOMA FLABELLARE | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA NIGRUM | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA SPECTABILE | 1 | 1 | 1 | 1 | 4 |
| FUNDULUS NOTATUS | 1 | 1 | 1 | 1 | 4 |
| GAMBUSIA AFFINIS | 1 | 1 | 1 | 1 | 4 |
| ICTALURUS PUNCTATUS | 0 | 0 | 1 | 1 | 2 |
| LEPISOSTEUS PLATOSTOMUS | 0 | 0 | 1 | 0 | 1 |
| LEPOMIS CYANELLUS | 0 | 0 | 0 | 0 | 0 |
| LEPOMIS HUMILIS | 0 | 1 | 1 | 1 | 3 |
| LEPOMIS MACROCHIRUS | 0 | 0 | 1 | 0 | 1 |
| LEPOMIS MEGALOTIS | 1 | 0 | 0 | 1 | 2 |
| LUXILUS CHRYSOCEPHALUS | 1 | 1 | 1 | 1 | 4 |
| LUXILUS CORNUTUS | 1 | 1 | 1 | 1 | 4 |
| LYTHRURUS UMBRATILIS | 1 | 1 | 0 | 1 | 3 |
| MICROPTERUS SALMOIDES | 0 | 0 | 1 | 1 | 2 |
| MOXOSTOMA ERYTHRURUM | 1 | 1 | 1 | 1 | 4 |
| NOTEMIGONUS CRYSOLEUCAS | 1 | 1 | 0 | 0 | 2 |
| NOTROPIS BOOPS | 1 | 1 | 1 | 0 | 3 |
| NOTROPIS DORSALIS | 1 | 1 | 1 | 1 | 4 |
| NOTROPIS STRAMINEUS | 1 | 1 | 1 | 1 | 4 |
| NOTURUS EXILIS | 1 | 1 | 1 | 1 | 4 |
| NOTURUS FLAVUS | 1 | 1 | 1 | 1 | 4 |
| PERCINA CAPRODES | 0 | 0 | 1 | 1 | 2 |
| PERCINA MACULATA | 1 | 0 | 0 | 0 | 1 |
| PERCINA PHOXOCEPHALA | 1 | 1 | 1 | 1 | 4 |

Plains Subregion

| Species | MARS | GAM | BRT | RndmFrst | Sum |
|-------------------------|------|-----|-----|----------|-----|
| PHENACOBIUS MIRABILIS | 1 | 0 | 1 | 0 | 2 |
| PIMEPHALES NOTATUS | 1 | 1 | 1 | 1 | 4 |
| PIMEPHALES PROMELAS | 1 | 1 | 1 | 1 | 4 |
| POMOXIS ANNULARIS | 0 | 1 | 1 | 1 | 3 |
| PYLODICTIS OLIVARIS | 1 | 0 | 1 | 0 | 2 |
| SEMOTILUS ATROMACULATUS | 1 | 1 | 1 | 1 | 4 |

Ozarks Subregion

| Species | MARS | BRT | GAM | RndmFrst | Sum |
|----------------------------|------|-----|-----|----------|-----|
| AMBLOPLITES ARIOMMUS | 0 | 1 | 0 | 1 | 2 |
| AMBLOPLITES CONSTELLATUS | 0 | 1 | 0 | 1 | 2 |
| AMBLOPLITES RUPESTRIS | 1 | 1 | 1 | 1 | 4 |
| AMEIURUS MELAS | 0 | 1 | 1 | 1 | 3 |
| AMEIURUS NATALIS | 0 | 0 | 0 | 1 | 1 |
| CAMPOSTOMA ANOMALUM PULLUM | 1 | 1 | 1 | 1 | 4 |
| CAMPOSTOMA OLIGOLEPIS | 1 | 1 | 1 | 1 | 4 |
| CATOSTOMUS COMMERSONII | 1 | 1 | 0 | 1 | 3 |
| COTTUS BAIRDII | 1 | 1 | 1 | 1 | 4 |
| COTTUS CAROLINAE | 1 | 1 | 1 | 1 | 4 |
| COTTUS HYPSELURUS | 1 | 1 | 1 | 1 | 4 |
| CYPRINELLA GALACTURA | 0 | 0 | 1 | 1 | 2 |
| CYPRINELLA LUTRENSIS | 1 | 1 | 1 | 1 | 4 |
| CYPRINELLA WHIPPLEI | 1 | 0 | 1 | 1 | 3 |
| DOROSOMA CEPEDIANUM | 0 | 1 | 0 | 0 | 1 |
| ERIMYZON OBLONGUS | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA BLENNIOIDES | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA CAERULEUM | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA FLABELLARE | 1 | 1 | 0 | 1 | 3 |
| ETHEOSTOMA NIGRUM | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA PUNCTULATUM | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA SPECTABILE | 1 | 1 | 1 | 1 | 4 |
| ETHEOSTOMA TETRAZONUM | 1 | 0 | 0 | 1 | 2 |
| ETHEOSTOMA ZONALE | 1 | 1 | 1 | 1 | 4 |
| FUNDULUS CATENATUS | 1 | 1 | 1 | 1 | 4 |
| FUNDULUS NOTATUS | 1 | 1 | 1 | 1 | 4 |
| FUNDULUS OLIVACEUS | 1 | 1 | 1 | 1 | 4 |
| FUNDULUS SCIADICUS | 1 | 0 | 1 | 1 | 3 |
| GAMBUSIA AFFINIS | 1 | 1 | 1 | 1 | 4 |
| HYBOPSIS AMBLOPS | 1 | 1 | 1 | 1 | 4 |
| HYPENTELIUM NIGRICANS | 0 | 1 | 1 | 1 | 3 |
| ICTALURUS PUNCTATUS | 0 | 1 | 0 | 1 | 2 |

| Species | MARS | BRT | GAM | RndmFrst | Sum |
|--------------------------|------|-----|-----|----------|-----|
| LEPISOSTEUS OSSEUS | 1 | 0 | 0 | 1 | 2 |
| LEPOMIS CYANELLUS | 0 | 1 | 1 | 1 | 3 |
| LEPOMIS GULOSUS | 1 | 1 | 1 | 0 | 3 |
| LEPOMIS HUMILIS | 1 | 1 | 1 | 1 | 4 |
| LEPOMIS MACROCHIRUS | 0 | 1 | 1 | 1 | 3 |
| LEPOMIS MEGALOTIS | 1 | 1 | 0 | 1 | 3 |
| LEPOMIS MICROLOPHUS | 0 | 0 | 1 | 1 | 2 |
| LUXILUS CARDINALIS | 1 | 1 | 1 | 1 | 4 |
| LUXILUS CHRYSOCEPHALUS | 1 | 1 | 1 | 1 | 4 |
| LUXILUS CORNUTUS | 1 | 1 | 1 | 1 | 4 |
| LUXILUS PILSBRYI | 1 | 1 | 1 | 1 | 4 |
| LUXILUS ZONATUS | 1 | 1 | 1 | 1 | 4 |
| LYTHRURUS UMBRATILIS | 0 | 1 | 1 | 1 | 3 |
| MICROPTERUS DOLOMIEU | 1 | 1 | 0 | 1 | 3 |
| MICROPTERUS PUNCTULATUS | 1 | 1 | 1 | 1 | 4 |
| MICROPTERUS SALMOIDES | 0 | 1 | 1 | 0 | 2 |
| MINYTREMA MELANOPS | 1 | 1 | 0 | 1 | 3 |
| MOXOSTOMA DUQUESNEI | 1 | 1 | 1 | 1 | 4 |
| MOXOSTOMA ERYTHRURUM | 1 | 1 | 1 | 1 | 4 |
| MOXOSTOMA MACROLEPIDOTUM | 1 | 0 | 1 | 1 | 3 |
| NOCOMIS ASPER | 0 | 1 | 1 | 1 | 3 |
| NOCOMIS BIGUTTATUS | 1 | 1 | 1 | 1 | 4 |
| NOTEMIGONUS CRYSOLEUCAS | 1 | 1 | 0 | 1 | 3 |
| NOTROPIS BOOPS | 1 | 1 | 1 | 1 | 4 |
| NOTROPIS GREENEI | 1 | 1 | 1 | 1 | 4 |
| NOTROPIS NUBILUS | 0 | 1 | 1 | 1 | 3 |
| NOTROPIS STRAMINEUS | 1 | 1 | 1 | 1 | 4 |
| NOTROPIS TELESCOPUS | 1 | 1 | 1 | 1 | 4 |
| NOTURUS ALBATER | 0 | 0 | 1 | 1 | 2 |
| NOTURUS EXILIS | 0 | 1 | 1 | 1 | 3 |
| PERCINA CAPRODES | 1 | 1 | 1 | 1 | 4 |
| PHENACOBIUS MIRABILIS | 1 | 1 | 1 | 1 | 4 |
| PHOXINUS ERYTHROGASTER | 1 | 1 | 1 | 1 | 4 |
| PIMEPHALES NOTATUS | 1 | 1 | 1 | 1 | 4 |
| PIMEPHALES PROMELAS | 0 | 1 | 0 | 0 | 1 |
| SEMOTILUS ATROMACULATUS | 1 | 1 | 1 | 1 | 4 |

Appendix 8: Component model evaluation statistics

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| AMEIURUS MELAS | 0.570 | 0.860 | 0.208 |
| AMEIURUS NATALIS | 0.597 | 1.055 | 0.139 |
| APLODINOTUS GRUNNIENS | 0.645 | 0.571 | 0.130 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.766 | 0.972 | 0.087 |
| CARPIODES CARPIO | 0.884 | 0.785 | 0.077 |
| CARPIODES CYPRINUS | 0.726 | 1.231 | 0.042 |
| CATOSTOMUS COMMERSONII | 0.667 | 1.120 | 0.084 |
| CYPRINELLA LUTRENSIS | 0.736 | 0.925 | 0.087 |
| DOROSOMA CEPEDIANUM | 0.846 | 1.092 | 0.051 |
| ETHEOSTOMA FLABELLARE | 0.908 | 0.906 | 0.064 |
| ETHEOSTOMA NIGRUM | 0.813 | 0.936 | 0.109 |
| ETHEOSTOMA SPECTABILE | 0.858 | 0.934 | 0.093 |
| FUNDULUS NOTATUS | 0.900 | 1.142 | 0.040 |
| GAMBUSIA AFFINIS | 0.759 | 1.024 | 0.083 |
| ICTALURUS PUNCTATUS | 0.816 | 0.795 | 0.128 |
| LEPISOSTEUS PLATOSTOMUS | 0.887 | 0.743 | 0.063 |
| LEPOMIS CYANELLUS | 0.517 | 0.951 | 0.146 |
| LEPOMIS HUMILIS | 0.699 | 0.810 | 0.113 |
| LEPOMIS MACROCHIRUS | 0.584 | 1.062 | 0.093 |
| LEPOMIS MEGALOTIS | 0.816 | 0.565 | 0.115 |
| LUXILUS CHRYSOCEPHALUS | 0.898 | 0.944 | 0.045 |
| LUXILUS CORNUTUS | 0.899 | 0.841 | 0.063 |
| LYTHRURUS UMBRATILIS | 0.711 | 0.941 | 0.099 |
| MICROPTERUS SALMOIDES | 0.575 | 0.967 | 0.116 |
| MOXOSTOMA ERYTHRURUM | 0.795 | 1.105 | 0.059 |
| NOTEMIGONUS CRYSOLEUCAS | 0.637 | 1.000 | 0.103 |
| NOTROPIS BOOPS | 0.671 | 1.004 | 0.082 |
| NOTROPIS DORSALIS | 0.877 | 0.963 | 0.061 |
| NOTROPIS STRAMINEUS | 0.722 | 0.947 | 0.091 |
| NOTURUS EXILIS | 0.930 | 0.795 | 0.076 |
| NOTURUS FLAVUS | 0.696 | 1.074 | 0.041 |
| PERCINA CAPRODES | 0.831 | 0.675 | 0.088 |
| PERCINA MACULATA | 0.918 | 0.700 | 0.091 |
| PERCINA PHOXOCEPHALA | 0.953 | 1.065 | 0.076 |
| PHENACOBIUS MIRABILIS | 0.672 | 0.950 | 0.143 |
| PIMEPHALES NOTATUS | 0.788 | 1.014 | 0.087 |
| PIMEPHALES PROMELAS | 0.755 | 0.961 | 0.112 |
| POMOXIS ANNULARIS | 0.791 | 0.833 | 0.099 |

Generalized Additive Models: Plains Subregion

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| PYLODICTIS OLIVARIS | 0.740 | 0.747 | 0.076 |
| SEMOTILUS ATROMACULATUS | 0.701 | 0.960 | 0.091 |

Generalized Additive Models: Ozark Subregion

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|------|
| AMBLOPLITES ARIOMMUS | 0.918 | 0.631 | 0.09 |
| AMBLOPLITES CONSTELLATUS | 0.682 | 1.778 | 0.02 |
| AMBLOPLITES RUPESTRIS | 0.892 | 1.012 | 0.05 |
| AMEIURUS MELAS | 0.714 | 0.990 | 0.04 |
| AMEIURUS NATALIS | 0.651 | 0.977 | 0.12 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.751 | 1.010 | 0.07 |
| CAMPOSTOMA OLIGOLEPIS | 0.836 | 0.949 | 0.07 |
| CATOSTOMUS COMMERSONII | 0.841 | 0.745 | 0.08 |
| COTTUS BAIRDII | 0.910 | 0.925 | 0.07 |
| COTTUS CAROLINAE | 0.829 | 0.986 | 0.07 |
| COTTUS HYPSELURUS | 0.884 | 0.816 | 0.07 |
| CYPRINELLA GALACTURA | 0.875 | 1.011 | 0.04 |
| CYPRINELLA LUTRENSIS | 0.940 | 0.914 | 0.07 |
| CYPRINELLA WHIPPLEI | 0.955 | 0.770 | 0.06 |
| DOROSOMA CEPEDIANUM | 0.628 | 1.536 | 0.06 |
| ERIMYZON OBLONGUS | 0.915 | 0.813 | 0.05 |
| ETHEOSTOMA BLENNIOIDES | 0.825 | 0.888 | 0.07 |
| ETHEOSTOMA CAERULEUM | 0.927 | 0.963 | 0.06 |
| ETHEOSTOMA FLABELLARE | 0.737 | 0.746 | 0.11 |
| ETHEOSTOMA NIGRUM | 0.872 | 0.840 | 0.06 |
| ETHEOSTOMA PUNCTULATUM | 0.874 | 0.962 | 0.05 |
| ETHEOSTOMA SPECTABILE | 0.813 | 0.933 | 0.06 |
| ETHEOSTOMA TETRAZONUM | 0.911 | 0.666 | 0.06 |
| ETHEOSTOMA ZONALE | 0.818 | 0.955 | 0.05 |
| FUNDULUS CATENATUS | 0.843 | 1.061 | 0.09 |
| FUNDULUS NOTATUS | 0.826 | 1.122 | 0.07 |
| FUNDULUS OLIVACEUS | 0.857 | 0.967 | 0.05 |
| FUNDULUS SCIADICUS | 0.955 | 0.935 | 0.05 |
| GAMBUSIA AFFINIS | 0.738 | 1.026 | 0.11 |
| HYBOPSIS AMBLOPS | 0.860 | 0.905 | 0.09 |
| HYPENTELIUM NIGRICANS | 0.721 | 1.043 | 0.10 |
| ICTALURUS PUNCTATUS | 0.750 | 1.324 | 0.04 |
| LEPISOSTEUS OSSEUS | 0.658 | 0.711 | 0.05 |
| LEPOMIS CYANELLUS | 0.646 | 0.974 | 0.09 |
| LEPOMIS GULOSUS | 0.761 | 0.895 | 0.05 |

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| LEPOMIS HUMILIS | 0.823 | 1.214 | 0.033 |
| LEPOMIS MACROCHIRUS | 0.673 | 1.036 | 0.088 |
| LEPOMIS MEGALOTIS | 0.645 | 0.985 | 0.126 |
| LEPOMIS MICROLOPHUS | 0.818 | 1.008 | 0.046 |
| LUXILUS CARDINALIS | 0.901 | 1.157 | 0.021 |
| LUXILUS CHRYSOCEPHALUS | 0.828 | 1.122 | 0.061 |
| LUXILUS CORNUTUS | 0.967 | 1.179 | 0.046 |
| LUXILUS PILSBRYI | 0.963 | 1.032 | 0.039 |
| LUXILUS ZONATUS | 0.921 | 0.984 | 0.037 |
| LYTHRURUS UMBRATILIS | 0.788 | 0.905 | 0.077 |
| MICROPTERUS DOLOMIEU | 0.766 | 1.001 | 0.128 |
| MICROPTERUS PUNCTULATUS | 0.818 | 0.993 | 0.060 |
| MICROPTERUS SALMOIDES | 0.696 | 1.003 | 0.101 |
| MINYTREMA MELANOPS | 0.875 | 1.849 | 0.031 |
| MOXOSTOMA DUQUESNEI | 0.752 | 1.002 | 0.089 |
| MOXOSTOMA ERYTHRURUM | 0.720 | 1.078 | 0.114 |
| MOXOSTOMA MACROLEPIDOTUM | 0.689 | 0.950 | 0.044 |
| NOCOMIS ASPER | 0.911 | 0.833 | 0.056 |
| NOCOMIS BIGUTTATUS | 0.851 | 0.947 | 0.109 |
| NOTEMIGONUS CRYSOLEUCAS | 0.671 | 0.750 | 0.070 |
| NOTROPIS BOOPS | 0.835 | 1.197 | 0.096 |
| NOTROPIS GREENEI | 0.920 | 0.997 | 0.078 |
| NOTROPIS NUBILUS | 0.762 | 0.977 | 0.076 |
| NOTROPIS STRAMINEUS | 0.874 | 1.175 | 0.058 |
| NOTROPIS TELESCOPUS | 0.944 | 0.938 | 0.081 |
| NOTURUS ALBATER | 0.939 | 0.964 | 0.102 |
| NOTURUS EXILIS | 0.673 | 0.922 | 0.113 |
| PERCINA CAPRODES | 0.788 | 0.953 | 0.052 |
| PHENACOBIUS MIRABILIS | 0.808 | 1.043 | 0.044 |
| PHOXINUS ERYTHROGASTER | 0.810 | 1.010 | 0.075 |
| PIMEPHALES NOTATUS | 0.878 | 0.964 | 0.067 |
| PIMEPHALES PROMELAS | 0.595 | 0.602 | 0.067 |
| SEMOTILUS ATROMACULATUS | 0.767 | 1.011 | 0.088 |

Multivariate Adaptive Regression Splines: Plains Subregion

| Species | AUC | Bias | MAE |
|-----------------------|-------|---------|-------|
| AMEIURUS MELAS | 0.527 | 0.865 | 0.143 |
| AMEIURUS NATALIS | 0.598 | 3 1.002 | 0.164 |
| APLODINOTUS GRUNNIENS | 0.688 | 1.733 | 0.080 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.680 | 1.020 | 0.161 |

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| CARPIODES CARPIO | 0.725 | 0.834 | 0.093 |
| CARPIODES CYPRINUS | 0.815 | 0.817 | 0.065 |
| CATOSTOMUS COMMERSONII | 0.601 | 1.111 | 0.159 |
| CYPRINELLA LUTRENSIS | 0.874 | 1.052 | 0.076 |
| DOROSOMA CEPEDIANUM | 0.763 | 0.737 | 0.067 |
| ETHEOSTOMA FLABELLARE | 0.866 | 1.006 | 0.056 |
| ETHEOSTOMA NIGRUM | 0.785 | 1.041 | 0.107 |
| ETHEOSTOMA SPECTABILE | 0.904 | 0.943 | 0.080 |
| FUNDULUS NOTATUS | 0.914 | 0.960 | 0.064 |
| GAMBUSIA AFFINIS | 0.786 | 1.130 | 0.090 |
| ICTALURUS PUNCTATUS | 0.716 | 0.995 | 0.130 |
| LEPISOSTEUS PLATOSTOMUS | 0.713 | 1.267 | 0.040 |
| LEPOMIS CYANELLUS | 0.582 | 1.001 | 0.044 |
| LEPOMIS HUMILIS | 0.683 | 0.710 | 0.107 |
| LEPOMIS MACROCHIRUS | 0.565 | 0.980 | 0.120 |
| LEPOMIS MEGALOTIS | 0.829 | 0.946 | 0.063 |
| LUXILUS CHRYSOCEPHALUS | 0.844 | 0.847 | 0.054 |
| LUXILUS CORNUTUS | 0.865 | 1.221 | 0.054 |
| LYTHRURUS UMBRATILIS | 0.735 | 1.080 | 0.091 |
| MICROPTERUS SALMOIDES | 0.520 | 0.974 | 0.186 |
| MOXOSTOMA ERYTHRURUM | 0.879 | 0.751 | 0.064 |
| NOTEMIGONUS CRYSOLEUCAS | 0.653 | 1.194 | 0.123 |
| NOTROPIS BOOPS | 0.942 | 0.858 | 0.059 |
| NOTROPIS DORSALIS | 0.862 | 1.091 | 0.098 |
| NOTROPIS STRAMINEUS | 0.737 | 1.046 | 0.125 |
| NOTURUS EXILIS | 0.923 | 0.841 | 0.067 |
| NOTURUS FLAVUS | 0.650 | 0.896 | 0.047 |
| PERCINA CAPRODES | 0.859 | 1.439 | 0.046 |
| PERCINA MACULATA | 0.847 | 0.778 | 0.066 |
| PERCINA PHOXOCEPHALA | 0.931 | 0.823 | 0.077 |
| PHENACOBIUS MIRABILIS | 0.742 | 1.043 | 0.091 |
| PIMEPHALES NOTATUS | 0.756 | 0.942 | 0.097 |
| PIMEPHALES PROMELAS | 0.767 | 0.992 | 0.109 |
| POMOXIS ANNULARIS | 0.655 | 1.256 | 0.076 |
| PYLODICTIS OLIVARIS | 0.865 | 0.847 | 0.057 |
| SEMOTILUS ATROMACULATUS | 0.756 | 0.997 | 0.065 |

Multivariate Adaptive Regression Splines: Ozark Subregion

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| AMBLOPLITES ARIOMMUS | 0.895 | 1.254 | 0.040 |
| AMBLOPLITES CONSTELLATUS | 0.968 | 0.712 | 0.082 |

| Species | AUC | Bias | MAE |
|------------------------|-------|-------|-------|
| AMBLOPLITES RUPESTRIS | 0.870 | 0.886 | 0.109 |
| AMEIURUS MELAS | 0.442 | 0.555 | 0.064 |
| AMEIURUS NATALIS | 0.560 | 1.039 | 0.199 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.739 | 1.050 | 0.055 |
| CAMPOSTOMA OLIGOLEPIS | 0.847 | 1.067 | 0.088 |
| CATOSTOMUS COMMERSONII | 0.829 | 0.892 | 0.072 |
| COTTUS BAIRDII | 0.908 | 0.966 | 0.065 |
| COTTUS CAROLINAE | 0.827 | 0.989 | 0.087 |
| COTTUS HYPSELURUS | 0.748 | 0.969 | 0.050 |
| CYPRINELLA GALACTURA | 0.898 | 0.515 | 0.072 |
| CYPRINELLA LUTRENSIS | 0.967 | 0.902 | 0.055 |
| CYPRINELLA WHIPPLEI | 0.961 | 0.960 | 0.064 |
| DOROSOMA CEPEDIANUM | 0.651 | 1.429 | 0.046 |
| ERIMYZON OBLONGUS | 0.820 | 1.175 | 0.037 |
| ETHEOSTOMA BLENNIOIDES | 0.756 | 0.931 | 0.111 |
| ETHEOSTOMA CAERULEUM | 0.882 | 0.996 | 0.079 |
| ETHEOSTOMA FLABELLARE | 0.710 | 0.912 | 0.077 |
| ETHEOSTOMA NIGRUM | 0.847 | 1.088 | 0.082 |
| ETHEOSTOMA PUNCTULATUM | 0.817 | 1.145 | 0.088 |
| ETHEOSTOMA SPECTABILE | 0.747 | 1.111 | 0.105 |
| ETHEOSTOMA TETRAZONUM | 0.800 | 0.815 | 0.053 |
| ETHEOSTOMA ZONALE | 0.795 | 1.184 | 0.076 |
| FUNDULUS CATENATUS | 0.860 | 1.061 | 0.101 |
| FUNDULUS NOTATUS | 0.809 | 1.114 | 0.063 |
| FUNDULUS OLIVACEUS | 0.773 | 1.040 | 0.102 |
| FUNDULUS SCIADICUS | 0.804 | 0.771 | 0.056 |
| GAMBUSIA AFFINIS | 0.734 | 0.997 | 0.084 |
| HYBOPSIS AMBLOPS | 0.855 | 0.961 | 0.055 |
| HYPENTELIUM NIGRICANS | 0.660 | 0.961 | 0.139 |
| ICTALURUS PUNCTATUS | 0.872 | 1.412 | 0.033 |
| LEPISOSTEUS OSSEUS | 0.678 | 0.852 | 0.050 |
| LEPOMIS CYANELLUS | 0.630 | 1.160 | 0.161 |
| LEPOMIS GULOSUS | 0.657 | 1.070 | 0.026 |
| LEPOMIS HUMILIS | 0.801 | 1.044 | 0.036 |
| LEPOMIS MACROCHIRUS | 0.628 | 0.935 | 0.138 |
| LEPOMIS MEGALOTIS | 0.639 | 0.978 | 0.087 |
| LEPOMIS MICROLOPHUS | 0.600 | 0.588 | 0.077 |
| LUXILUS CARDINALIS | 0.993 | 0.873 | 0.053 |
| LUXILUS CHRYSOCEPHALUS | 0.852 | 1.066 | 0.067 |
| LUXILUS CORNUTUS | 0.926 | 1.007 | 0.053 |
| LUXILUS PILSBRYI | 0.996 | 0.961 | 0.046 |

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| LUXILUS ZONATUS | 0.913 | 1.015 | 0.049 |
| LYTHRURUS UMBRATILIS | 0.728 | 1.310 | 0.043 |
| MICROPTERUS DOLOMIEU | 0.749 | 1.051 | 0.108 |
| MICROPTERUS PUNCTULATUS | 0.826 | 0.930 | 0.049 |
| MICROPTERUS SALMOIDES | 0.572 | 1.084 | 0.179 |
| MINYTREMA MELANOPS | 0.769 | 1.083 | 0.038 |
| MOXOSTOMA DUQUESNEI | 0.717 | 0.931 | 0.109 |
| MOXOSTOMA ERYTHRURUM | 0.691 | 1.068 | 0.115 |
| MOXOSTOMA MACROLEPIDOTUM | 0.815 | 0.790 | 0.054 |
| NOCOMIS ASPER | 0.971 | 0.666 | 0.087 |
| NOCOMIS BIGUTTATUS | 0.900 | 1.033 | 0.066 |
| NOTEMIGONUS CRYSOLEUCAS | 0.654 | 0.965 | 0.033 |
| NOTROPIS BOOPS | 0.836 | 0.875 | 0.107 |
| NOTROPIS GREENEI | 0.905 | 1.248 | 0.042 |
| NOTROPIS NUBILUS | 0.758 | 1.084 | 0.126 |
| NOTROPIS STRAMINEUS | 0.912 | 1.069 | 0.056 |
| NOTROPIS TELESCOPUS | 0.950 | 0.919 | 0.094 |
| NOTURUS ALBATER | 0.894 | 1.328 | 0.061 |
| NOTURUS EXILIS | 0.665 | 1.264 | 0.143 |
| PERCINA CAPRODES | 0.661 | 1.209 | 0.084 |
| PHENACOBIUS MIRABILIS | 0.691 | 1.066 | 0.039 |
| PHOXINUS ERYTHROGASTER | 0.812 | 1.090 | 0.065 |
| PIMEPHALES NOTATUS | 0.845 | 0.935 | 0.088 |
| PIMEPHALES PROMELAS | 0.625 | 0.611 | 0.054 |
| SEMOTILUS ATROMACULATUS | 0.779 | 1.052 | 0.097 |

Boosted Regression Trees: Plains Subregion

| Species | AUC | Bias | MAE |
|------------------------|-------|-------|-------|
| AMEIURUS MELAS | 0.478 | 0.899 | 0.174 |
| AMEIURUS NATALIS | 0.684 | 0.858 | 0.116 |
| APLODINOTUS GRUNNIENS | 0.910 | 1.363 | 0.055 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.796 | 1.085 | 0.090 |
| CARPIODES CARPIO | 0.824 | 0.887 | 0.064 |
| CARPIODES CYPRINUS | 0.814 | 1.916 | 0.049 |
| CATOSTOMUS COMMERSONII | 0.724 | 1.036 | 0.126 |
| CYPRINELLA LUTRENSIS | 0.840 | 0.977 | 0.059 |
| DOROSOMA CEPEDIANUM | 0.803 | 1.450 | 0.072 |
| ETHEOSTOMA FLABELLARE | 0.943 | 1.125 | 0.055 |
| ETHEOSTOMA NIGRUM | 0.796 | 0.926 | 0.125 |
| ETHEOSTOMA SPECTABILE | 0.882 | 0.843 | 0.110 |

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| FUNDULUS NOTATUS | 0.969 | 1.166 | 0.082 |
| GAMBUSIA AFFINIS | 0.817 | 0.928 | 0.082 |
| ICTALURUS PUNCTATUS | 0.849 | 0.884 | 0.078 |
| LEPISOSTEUS PLATOSTOMUS | 0.845 | 0.835 | 0.053 |
| LEPOMIS HUMILIS | 0.711 | 1.042 | 0.064 |
| LEPOMIS MACROCHIRUS | 0.643 | 0.947 | 0.091 |
| LEPOMIS MEGALOTIS | 0.966 | 0.742 | 0.077 |
| LUXILUS CHRYSOCEPHALUS | 0.988 | 0.980 | 0.068 |
| LUXILUS CORNUTUS | 0.888 | 1.036 | 0.048 |
| LYTHRURUS UMBRATILIS | 0.715 | 0.892 | 0.136 |
| MICROPTERUS SALMOIDES | 0.647 | 1.017 | 0.095 |
| MOXOSTOMA ERYTHRURUM | 0.870 | 1.217 | 0.041 |
| NOTEMIGONUS CRYSOLEUCAS | 0.643 | 0.817 | 0.135 |
| NOTROPIS BOOPS | 0.964 | 0.760 | 0.086 |
| NOTROPIS DORSALIS | 0.897 | 1.038 | 0.072 |
| NOTROPIS STRAMINEUS | 0.792 | 0.870 | 0.119 |
| NOTURUS EXILIS | 0.952 | 1.031 | 0.073 |
| NOTURUS FLAVUS | 0.702 | 1.054 | 0.045 |
| PERCINA CAPRODES | 0.865 | 1.034 | 0.075 |
| PERCINA MACULATA | 0.864 | 2.491 | 0.046 |
| PERCINA PHOXOCEPHALA | 0.947 | 1.150 | 0.061 |
| PHENACOBIUS MIRABILIS | 0.666 | 1.104 | 0.112 |
| PIMEPHALES NOTATUS | 0.823 | 1.021 | 0.033 |
| PIMEPHALES PROMELAS | 0.847 | 0.880 | 0.082 |
| POMOXIS ANNULARIS | 0.676 | 1.071 | 0.092 |
| PYLODICTIS OLIVARIS | 0.917 | 0.880 | 0.063 |
| SEMOTILUS ATROMACULATUS | 0.811 | 0.999 | 0.057 |

Boosted Regression Trees: Ozark Subregion

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| AMBLOPLITES ARIOMMUS | 0.916 | 0.924 | 0.080 |
| AMBLOPLITES CONSTELLATUS | 0.979 | 0.779 | 0.068 |
| AMBLOPLITES RUPESTRIS | 0.909 | 0.911 | 0.057 |
| AMEIURUS MELAS | 0.759 | 0.818 | 0.041 |
| AMEIURUS NATALIS | 0.693 | 1.203 | 0.128 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.827 | 1.011 | 0.030 |
| CAMPOSTOMA OLIGOLEPIS | 0.891 | 1.010 | 0.094 |
| CATOSTOMUS COMMERSONII | 0.826 | 0.925 | 0.053 |
| COTTUS BAIRDII | 0.924 | 1.146 | 0.047 |
| COTTUS CAROLINAE | 0.824 | 0.929 | 0.088 |

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| COTTUS HYPSELURUS | 0.852 | 0.877 | 0.072 |
| CYPRINELLA GALACTURA | 0.912 | 2.094 | 0.030 |
| CYPRINELLA LUTRENSIS | 0.942 | 0.902 | 0.070 |
| CYPRINELLA WHIPPLEI | 0.977 | 1.412 | 0.066 |
| DOROSOMA CEPEDIANUM | 0.739 | 0.908 | 0.033 |
| ERIMYZON OBLONGUS | 0.784 | 0.889 | 0.073 |
| ETHEOSTOMA BLENNIOIDES | 0.819 | 0.991 | 0.066 |
| ETHEOSTOMA CAERULEUM | 0.920 | 0.985 | 0.063 |
| ETHEOSTOMA FLABELLARE | 0.743 | 0.907 | 0.095 |
| ETHEOSTOMA NIGRUM | 0.854 | 0.981 | 0.052 |
| ETHEOSTOMA PUNCTULATUM | 0.871 | 0.928 | 0.067 |
| ETHEOSTOMA SPECTABILE | 0.827 | 1.040 | 0.059 |
| ETHEOSTOMA TETRAZONUM | 0.927 | 1.528 | 0.056 |
| ETHEOSTOMA ZONALE | 0.859 | 1.095 | 0.061 |
| FUNDULUS CATENATUS | 0.887 | 1.034 | 0.061 |
| FUNDULUS NOTATUS | 0.835 | 0.957 | 0.074 |
| FUNDULUS OLIVACEUS | 0.831 | 1.030 | 0.067 |
| FUNDULUS SCIADICUS | 0.922 | 1.396 | 0.044 |
| GAMBUSIA AFFINIS | 0.786 | 0.927 | 0.059 |
| HYBOPSIS AMBLOPS | 0.907 | 0.806 | 0.060 |
| HYPENTELIUM NIGRICANS | 0.733 | 0.926 | 0.087 |
| ICTALURUS PUNCTATUS | 0.872 | 1.129 | 0.046 |
| LEPISOSTEUS OSSEUS | 0.624 | 1.406 | 0.027 |
| LEPOMIS CYANELLUS | 0.687 | 0.969 | 0.080 |
| LEPOMIS GULOSUS | 0.697 | 1.020 | 0.031 |
| LEPOMIS HUMILIS | 0.907 | 1.144 | 0.026 |
| LEPOMIS MACROCHIRUS | 0.698 | 1.032 | 0.092 |
| LEPOMIS MEGALOTIS | 0.691 | 0.987 | 0.115 |
| LEPOMIS MICROLOPHUS | 0.838 | 1.367 | 0.072 |
| LUXILUS CARDINALIS | 0.999 | 1.117 | 0.046 |
| LUXILUS CHRYSOCEPHALUS | 0.872 | 1.013 | 0.038 |
| LUXILUS CORNUTUS | 0.979 | 1.038 | 0.057 |
| LUXILUS PILSBRYI | 1.000 | 1.033 | 0.051 |
| LUXILUS ZONATUS | 0.961 | 0.981 | 0.059 |
| LYTHRURUS UMBRATILIS | 0.830 | 1.061 | 0.038 |
| MICROPTERUS DOLOMIEU | 0.833 | 0.946 | 0.080 |
| MICROPTERUS PUNCTULATUS | 0.767 | 1.225 | 0.059 |
| MICROPTERUS SALMOIDES | 0.732 | 0.975 | 0.079 |
| MINYTREMA MELANOPS | 0.797 | 0.841 | 0.046 |
| MOXOSTOMA DUQUESNEI | 0.784 | 1.049 | 0.046 |
| MOXOSTOMA ERYTHRURUM | 0.793 | 1.052 | 0.046 |
| MOXOSTOMA MACROLEPIDOTUM | 0.733 | 1.480 | 0.037 |

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| NOCOMIS ASPER | 0.988 | 0.933 | 0.070 |
| NOCOMIS BIGUTTATUS | 0.878 | 0.944 | 0.085 |
| NOTEMIGONUS CRYSOLEUCAS | 0.637 | 1.076 | 0.072 |
| NOTROPIS BOOPS | 0.846 | 0.982 | 0.052 |
| NOTROPIS GREENEI | 0.934 | 1.132 | 0.064 |
| NOTROPIS NUBILUS | 0.827 | 1.000 | 0.085 |
| NOTROPIS STRAMINEUS | 0.882 | 0.974 | 0.091 |
| NOTROPIS TELESCOPUS | 0.944 | 0.820 | 0.082 |
| NOTURUS ALBATER | 0.963 | 1.489 | 0.067 |
| NOTURUS EXILIS | 0.703 | 0.870 | 0.105 |
| PERCINA CAPRODES | 0.804 | 0.924 | 0.038 |
| PHENACOBIUS MIRABILIS | 0.858 | 1.018 | 0.060 |
| PHOXINUS ERYTHROGASTER | 0.838 | 0.933 | 0.065 |
| PIMEPHALES NOTATUS | 0.882 | 0.984 | 0.058 |
| PIMEPHALES PROMELAS | 0.607 | 0.920 | 0.083 |
| SEMOTILUS ATROMACULATUS | 0.807 | 0.944 | 0.082 |

Random Forest: Plains Subregion

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| AMEIURUS MELAS | 0.697 | 0.746 | 0.240 |
| AMEIURUS NATALIS | 0.606 | 0.950 | 0.117 |
| APLODINOTUS GRUNNIENS | 0.949 | 1.137 | 0.073 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.842 | 0.925 | 0.078 |
| CARPIODES CARPIO | 0.779 | 1.088 | 0.054 |
| CARPIODES CYPRINUS | 0.897 | 0.829 | 0.112 |
| CATOSTOMUS COMMERSONII | 0.692 | 1.059 | 0.097 |
| CYPRINELLA LUTRENSIS | 0.860 | 1.015 | 0.072 |
| DOROSOMA CEPEDIANUM | 0.796 | 1.397 | 0.042 |
| ETHEOSTOMA FLABELLARE | 0.905 | 0.817 | 0.069 |
| ETHEOSTOMA NIGRUM | 0.855 | 1.027 | 0.077 |
| ETHEOSTOMA SPECTABILE | 0.928 | 0.818 | 0.124 |
| FUNDULUS NOTATUS | 0.940 | 1.029 | 0.059 |
| GAMBUSIA AFFINIS | 0.761 | 0.868 | 0.106 |
| ICTALURUS PUNCTATUS | 0.891 | 1.042 | 0.078 |
| LEPISOSTEUS PLATOSTOMUS | 0.784 | 0.738 | 0.115 |
| LEPOMIS CYANELLUS | 0.542 | 0.959 | 0.134 |
| LEPOMIS HUMILIS | 0.770 | 1.313 | 0.091 |
| LEPOMIS MACROCHIRUS | 0.527 | 1.110 | 0.189 |
| LEPOMIS MEGALOTIS | 0.943 | 1.052 | 0.048 |
| LUXILUS CHRYSOCEPHALUS | 0.953 | 1.182 | 0.090 |

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| LUXILUS CORNUTUS | 0.964 | 1.048 | 0.077 |
| LYTHRURUS UMBRATILIS | 0.766 | 0.942 | 0.088 |
| MICROPTERUS SALMOIDES | 0.666 | 0.916 | 0.115 |
| MOXOSTOMA ERYTHRURUM | 0.910 | 1.167 | 0.049 |
| NOTEMIGONUS CRYSOLEUCAS | 0.545 | 0.971 | 0.181 |
| NOTROPIS BOOPS | 0.970 | 0.665 | 0.083 |
| NOTROPIS DORSALIS | 0.933 | 1.038 | 0.067 |
| NOTROPIS STRAMINEUS | 0.847 | 0.964 | 0.052 |
| NOTURUS EXILIS | 0.921 | 0.912 | 0.060 |
| NOTURUS FLAVUS | 0.728 | 0.763 | 0.079 |
| PERCINA CAPRODES | 0.774 | 1.588 | 0.078 |
| PERCINA MACULATA | 0.915 | 0.605 | 0.092 |
| PERCINA PHOXOCEPHALA | 0.909 | 1.138 | 0.043 |
| PHENACOBIUS MIRABILIS | 0.623 | 0.986 | 0.184 |
| PIMEPHALES NOTATUS | 0.822 | 1.121 | 0.111 |
| PIMEPHALES PROMELAS | 0.791 | 1.082 | 0.074 |
| POMOXIS ANNULARIS | 0.702 | 0.777 | 0.103 |
| PYLODICTIS OLIVARIS | 0.937 | 0.685 | 0.107 |
| SEMOTILUS ATROMACULATUS | 0.770 | 1.021 | 0.082 |

Random Forest: Ozark Subregion

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| AMBLOPLITES ARIOMMUS | 0.931 | 1.070 | 0.065 |
| AMBLOPLITES CONSTELLATUS | 0.982 | 1.075 | 0.064 |
| AMBLOPLITES RUPESTRIS | 0.931 | 0.931 | 0.057 |
| AMEIURUS MELAS | 0.673 | 1.237 | 0.070 |
| AMEIURUS NATALIS | 0.719 | 0.928 | 0.090 |
| CAMPOSTOMA ANOMALUM | | | |
| PULLUM | 0.783 | 0.982 | 0.073 |
| CAMPOSTOMA OLIGOLEPIS | 0.882 | 1.083 | 0.061 |
| CATOSTOMUS COMMERSONII | 0.847 | 0.986 | 0.051 |
| COTTUS BAIRDII | 0.917 | 1.068 | 0.039 |
| COTTUS CAROLINAE | 0.853 | 0.984 | 0.074 |
| COTTUS HYPSELURUS | 0.850 | 0.764 | 0.112 |
| CYPRINELLA GALACTURA | 0.909 | 0.789 | 0.062 |
| CYPRINELLA LUTRENSIS | 0.962 | 1.056 | 0.061 |
| CYPRINELLA WHIPPLEI | 0.973 | 0.788 | 0.063 |
| DOROSOMA CEPEDIANUM | 0.818 | 0.695 | 0.067 |
| ERIMYZON OBLONGUS | 0.824 | 0.791 | 0.098 |
| ETHEOSTOMA BLENNIOIDES | 0.874 | 0.980 | 0.076 |
| ETHEOSTOMA CAERULEUM | 0.924 | 1.011 | 0.051 |

| Species | AUC | Bias | MAE |
|--------------------------|-------|-------|-------|
| ETHEOSTOMA FLABELLARE | 0.813 | 1.042 | 0.065 |
| ETHEOSTOMA NIGRUM | 0.877 | 1.111 | 0.065 |
| ETHEOSTOMA PUNCTULATUM | 0.907 | 1.030 | 0.054 |
| ETHEOSTOMA SPECTABILE | 0.807 | 1.000 | 0.056 |
| ETHEOSTOMA TETRAZONUM | 0.964 | 0.999 | 0.067 |
| ETHEOSTOMA ZONALE | 0.864 | 0.874 | 0.063 |
| FUNDULUS CATENATUS | 0.860 | 0.996 | 0.060 |
| FUNDULUS NOTATUS | 0.868 | 1.136 | 0.036 |
| FUNDULUS OLIVACEUS | 0.880 | 1.033 | 0.071 |
| FUNDULUS SCIADICUS | 0.952 | 0.988 | 0.064 |
| GAMBUSIA AFFINIS | 0.755 | 0.932 | 0.080 |
| HYBOPSIS AMBLOPS | 0.890 | 1.240 | 0.054 |
| HYPENTELIUM NIGRICANS | 0.711 | 1.043 | 0.085 |
| ICTALURUS PUNCTATUS | 0.884 | 0.765 | 0.059 |
| LEPISOSTEUS OSSEUS | 0.753 | 0.905 | 0.063 |
| LEPOMIS CYANELLUS | 0.654 | 1.002 | 0.096 |
| LEPOMIS GULOSUS | 0.721 | 1.305 | 0.059 |
| LEPOMIS HUMILIS | 0.839 | 1.202 | 0.042 |
| LEPOMIS MACROCHIRUS | 0.720 | 1.046 | 0.123 |
| LEPOMIS MEGALOTIS | 0.720 | 0.983 | 0.108 |
| LEPOMIS MICROLOPHUS | 0.772 | 0.900 | 0.061 |
| LUXILUS CARDINALIS | 0.997 | 0.929 | 0.054 |
| LUXILUS CHRYSOCEPHALUS | 0.883 | 0.991 | 0.061 |
| LUXILUS CORNUTUS | 0.971 | 0.823 | 0.065 |
| LUXILUS PILSBRYI | 1.000 | 1.033 | 0.056 |
| LUXILUS ZONATUS | 0.950 | 0.929 | 0.071 |
| LYTHRURUS UMBRATILIS | 0.896 | 1.052 | 0.052 |
| MICROPTERUS DOLOMIEU | 0.848 | 1.005 | 0.078 |
| MICROPTERUS PUNCTULATUS | 0.786 | 1.209 | 0.035 |
| MICROPTERUS SALMOIDES | 0.638 | 0.945 | 0.138 |
| MINYTREMA MELANOPS | 0.845 | 0.814 | 0.061 |
| MOXOSTOMA DUQUESNEI | 0.762 | 0.984 | 0.076 |
| MOXOSTOMA ERYTHRURUM | 0.811 | 1.084 | 0.051 |
| MOXOSTOMA MACROLEPIDOTUM | 0.837 | 1.094 | 0.099 |
| NOCOMIS ASPER | 0.996 | 0.988 | 0.059 |
| NOCOMIS BIGUTTATUS | 0.916 | 0.881 | 0.083 |
| NOTEMIGONUS CRYSOLEUCAS | 0.654 | 0.817 | 0.102 |
| NOTROPIS BOOPS | 0.896 | 0.836 | 0.074 |
| NOTROPIS GREENEI | 0.914 | 0.917 | 0.057 |
| NOTROPIS NUBILUS | 0.827 | 1.041 | 0.065 |
| NOTROPIS STRAMINEUS | 0.937 | 0.990 | 0.068 |
| NOTROPIS TELESCOPUS | 0.965 | 0.858 | 0.065 |

| Species | AUC | Bias | MAE |
|-------------------------|-------|-------|-------|
| NOTURUS ALBATER | 0.918 | 1.146 | 0.053 |
| NOTURUS EXILIS | 0.700 | 0.916 | 0.106 |
| PERCINA CAPRODES | 0.775 | 0.980 | 0.043 |
| PHENACOBIUS MIRABILIS | 0.773 | 1.213 | 0.044 |
| PHOXINUS ERYTHROGASTER | 0.853 | 0.979 | 0.096 |
| PIMEPHALES NOTATUS | 0.873 | 1.055 | 0.065 |
| PIMEPHALES PROMELAS | 0.675 | 1.356 | 0.040 |
| SEMOTILUS ATROMACULATUS | 0.794 | 1.134 | 0.088 |

Appendix 9: Electronic data with PDF of species distribution models and a shapefile of species distribution model results.

| A_ARIOMMUS 333.224 A_CONSTELLATUS 191.073 A_GRUNNIENS 312.024 A_MELAS 356.84 A_NATALIS 1266.47 A_NEBULOSUS 12.516 A_RUPESTRIS 324.638 A_SAYANUS 77.158 C_CAMURA 27.348 C_CAROLINAE 832.542 C_COMMERSONII 548.134 C_CYPRINUS 198.531 C_GALACTURA 222.156 C HYPSELURUS 412.178 | 312.602 202.543 346.505 425.111 1371.009 10.38 407.552 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 190.134 370.077 276.972 680.19 1945.802 17.072 608.175 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 300.444 |
|--|--|--|
| A_GRUNNIENS312.024A_MELAS356.84A_NATALIS1266.47A_NEBULOSUS12.516A_RUPESTRIS324.638A_SAYANUS77.158C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 346.505 425.111 1371.009 10.38 407.552 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 276.972 680.19 1945.802 17.072 608.175 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| A_MELAS356.84A_NATALIS1266.47A_NEBULOSUS12.516A_RUPESTRIS324.638A_SAYANUS77.158C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 425.111 1371.009 10.38 407.552 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 680.19 1945.802 17.072 608.175 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| A_NATALIS1266.47A_NEBULOSUS12.516A_RUPESTRIS324.638A_SAYANUS77.158C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 1371.009 10.38 407.552 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 1945.802 17.072 608.175 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| A_NEBULOSUS12.516A_RUPESTRIS324.638A_SAYANUS77.158C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 10.38 407.552 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 17.072 608.175 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| A_RUPESTRIS324.638A_SAYANUS77.158C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 407.552 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 608.175 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| A_SAYANUS77.158C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 74.228 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 125.509 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| C_BAIRDII439.734C_CAMURA27.348C_CAROLINAE832.542C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 492.169 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 625.104 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| C_CAMURA 27.348 C_CAROLINAE 832.542 C_CARPIO 242.071 C_COMMERSONII 548.134 C_CYPRINUS 198.531 C_GALACTURA 222.156 | 6.978 930.886 384.908 742.002 254.096 190.053 464.614 | 13.322 1009.521 360.436 1282.065 329.692 229.144 |
| C_CAROLINAE832.542C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 930.886 384.908 742.002 254.096 190.053 464.614 | 1009.521 360.436 1282.065 329.692 229.144 |
| C_CARPIO242.071C_COMMERSONII548.134C_CYPRINUS198.531C_GALACTURA222.156 | 384.908 742.002 254.096 190.053 464.614 | 360.436 1282.065 329.692 229.144 |
| C_COMMERSONII 548.134 C_CYPRINUS 198.531 C_GALACTURA 222.156 | 742.002 254.096 190.053 464.614 | 1282.065 329.692 229.144 |
| C_CYPRINUS 198.531 C_GALACTURA 222.156 | 254.096 190.053 464.614 | 329.692 229.144 |
| C_GALACTURA 222.156 | 190.053 464.614 | 229.144 |
| — | 464.614 | |
| | | 300.444 |
| C_HTPSELURUS 412.176 | | |
| C_LUTRENSIS 912.407 | 1326.801 | 1810.257 |
| C_MACROPTERUS 33.6 | 15.8 | 6.2 |
| C_OLIGOLEPIS 1456.787 | 1507.307 | 1630.861 |
| C_PULLUM 2125.527 | 2330.205 | 3049.566 |
| C_SPILOPTERA 84.163 | 80.43 | 186.09 |
| C_VELIFER 6.104 | 8.204 | 11.708 |
| C_VENUSTA 170.177 | 146.401 | 208.932 |
| C_WHIPPLEI 158.337 | 187.606 | 491.829 |
| D_CEPEDIANUM 567.991 | 560.338 | 535.041 |
| E_BLENNIOIDES 1138.156 | 1146.546 | 1648.462 |
| E_BURRI 92.391 | 88.268 | 27.531 |
| E_CAERULEUM 1721.378 | 1720.852 | 2072.02 |
| E_CRAGINI 50.004 | 19.332 | 30.192 |
| E_EUZONUM 22.25 | 20.93 | 8.06 |
| E_FLABELLARE 918.181 | 926.81 | 1358.272 |
| E_GRACILE 58.168 | 54.488 | 55.796 |
| E_HARRYI 139.302 | 130.406 | 81.804 |
| E_JULIAE 95.721 | 107.23 | 173.854 |
| E_MICROPERCA 57.04 | 75.769 | 37.193 |
| E_NIANGUAE 33.792 | 37.879 | 54.502 |
| E_NIGER 106.802 | 83.763 | 57.2 |
| E_NIGRUM 564.169 | 790.43 | 1528.231 |
| E_OBLONGUS 299.748 | 264.634 | 398.976 |

Appendix 10: Number of stream segments within each established conservation network each species is predicted to occupy

| E PROELIARE | 55.124 | 58.456 | 58.104 |
|------------------|----------|----------|----------|
| E PUNCTULATUM | 427.052 | 493.718 | 725.927 |
| E SPECTABILE | 1176.104 | 1206.013 | 1931.835 |
| E STIGMAEUM | 86.117 | 73.076 | 58.28 |
| E_TETRAZONUM | 183.452 | 236.027 | 260.582 |
| E UNIPORUM | 180.541 | 168.859 | 54.685 |
| E VERMICULATUS | 125.286 | 115.482 | 110.629 |
| E XPUNCTATUS | 57.672 | 63.74 | 132.266 |
| E ZONALE | 476.071 | 481.235 | 570.76 |
| F CATENATUS | 1557.236 | 1549.964 | 1737.935 |
| F DISPAR | 31.354 | 12.418 | 10.51 |
| F NOTATUS | 600.436 | 576.15 | 1139.801 |
| F OLIVACEUS | 1449.142 | 1433.267 | 2005.976 |
| F SCIADICUS | 105.019 | 146.294 | 111.938 |
| G AFFINIS | 988.319 | 1123.657 | 1391.275 |
| H_AMBLOPS | 489.888 | 398.22 | 607.802 |
| H_ARGYRITIS | 3.842 | 7.751 | 11.333 |
| H_HANKINSONI | 34.222 | 54.862 | 106.378 |
| H_NIGRICANS | 1317.83 | 1316.261 | 1632.585 |
| H_PLACITUS | 7.312 | 13.952 | 21.573 |
| I_BUBALUS | 166.85 | 125.569 | 156.312 |
| I_CYPRINELLUS | 143.521 | 115.563 | 129.897 |
| I_FOSSOR | 5.642 | 5.385 | 16.537 |
| I_GAGEI | 5.48 | 5.152 | 5.376 |
| I_NIGER | 2.96 | 5.288 | 7.272 |
| I_PUNCTATUS | 577.307 | 785.867 | 845.382 |
| L_AEPYPTERA | 102.171 | 99.879 | 59.139 |
| L_CARDINALIS | 96.268 | 87.251 | 147.858 |
| L_CHRYSOCEPHALUS | 1167.124 | 1173.394 | 1922.021 |
| L_CORNUTUS | 179.518 | 225.801 | 337.47 |
| L_CYANELLUS | 2397.419 | 2709.253 | 3684.798 |
| L_GULOSUS | 345.131 | 289.807 | 271.16 |
| L_HUMILIS | 469.552 | 482.951 | 739.338 |
| L_MACROCHIRUS | 2418.909 | 2615.015 | 3471.637 |
| L_MEGALOTIS | 2064.565 | 1970.876 | 2438.593 |
| L_MICROLOPHUS | 185.973 | 175.574 | 282.198 |
| L_MINIATUS | 196.628 | 194.65 | 163.016 |
| L_OCULATUS | 114.438 | 74.358 | 149.09 |
| L_OSSEUS | 219.758 | 185.294 | 236.645 |
| L_PILSBRYI | 354.921 | 383.906 | 607.76 |
| L_PLATOSTOMUS | 337.486 | 345.116 | 234.484 |
| L_UMBRATILIS | 489.611 | 538.704 | 1079.536 |
| L_ZONATUS | 1566.112 | 1579.114 | 1625.662 |
| | | | |

| M ANISURUM | 48.501 | 59.233 | 122.735 |
|------------------|----------|----------|----------|
| M CARINATUM | 14.184 | 20.616 | 41.304 |
| M CHRYSOPS | 53.952 | 59.586 | 51.379 |
| M DOLOMIEU | 1197.13 | 1232.507 | 1469.864 |
| M DUQUESNEI | 567.748 | 614.134 | 857.632 |
| M ERYTHRURUM | 658.646 | 649.264 | 1059.4 |
| M_MACROLEPIDOTUM | 190.842 | 193.195 | 237.92 |
| M_MELANOPS | 225.481 | 145.166 | 177.67 |
| M_PUNCTULATUS | 400.249 | 383.568 | 472.881 |
| M_SALMOIDES | 1664.5 | 1857.451 | 2758.534 |
| M_STORERIANA | 15.78 | 19.042 | 24.607 |
| N_ALBATER | 413.477 | 358.548 | 260.499 |
| N_ASPER | 42.447 | 50.022 | 94.257 |
| N_ATHERINOIDES | 219.081 | 185.473 | 259.305 |
| N_BIGUTTATUS | 1388.826 | 1380.565 | 1458.036 |
| N_BOOPS | 751.893 | 681.436 | 1267.109 |
| N_BUCCATUS | 57.893 | 57.814 | 119.696 |
| N_BUCHANANI | 22.862 | 40.274 | 51.512 |
| N_CRYSOLEUCAS | 477.946 | 574.028 | 778.319 |
| N_DORSALIS | 313.691 | 638.961 | 1139.174 |
| N_EXILIS | 1230.81 | 1282.544 | 1899.379 |
| N_FLAVATER | 68.018 | 59.236 | 41.846 |
| N_FLAVUS | 119.489 | 182.883 | 238.85 |
| N_GREENEI | 494.088 | 476.583 | 463.179 |
| N_GYRINUS | 80.633 | 109.355 | 206.751 |
| N_HETEROLEPIS | 33.93 | 42.708 | 26.84 |
| N_MIURUS | 79.458 | 69.785 | 29.361 |
| N_NOCTURNUS | 49.718 | 47.006 | 36.682 |
| N_NUBILUS | 1536.215 | 1514.813 | 1737.241 |
| N_OZARCANUS | 119.198 | 110.086 | 83.054 |
| N_STRAMINEUS | 670.381 | 1117.532 | 1693.882 |
| N_TELESCOPUS | 683.748 | 609.556 | 516.686 |
| N_TEXANUS | 57.732 | 59.286 | 104.15 |
| N_VOLUCELLUS | 221.653 | 152.981 | 286.023 |
| O_EMILIAE | 87.132 | 76.933 | 84.607 |
| P_ANNULARIS | 351.843 | 452.291 | 441.372 |
| P_CAPRODES | 491.314 | 453.792 | 744.809 |
| P_COPELANDI | 20.935 | 4.969 | 8.262 |
| P_CYMATOTAENIA | 49.108 | 76.411 | 30.437 |
| P_ERYTHROGASTER | 798.875 | 849.866 | 908.488 |
| P_EVIDES | 87.196 | 76.751 | 132.765 |
| P_FLAVESCENS | 3.222 | 3.573 | 5.724 |
| P_MACULATA | 85.65 | 160.374 | 284.123 |

| P_MIRABILIS | 422.573 | 685.439 | 989.015 |
|------------------|----------|----------|----------|
| P_NIGROMACULATUS | 117.195 | 123.195 | 167.549 |
| P_NOTATUS | 1653.102 | 1948.989 | 2984.05 |
| P_OLIVARIS | 209.695 | 264.393 | 209.786 |
| P_PHOXOCEPHALA | 164.651 | 228.351 | 198.93 |
| P_PROMELAS | 417.229 | 694.736 | 1116.33 |
| P_SCIERA | 48.423 | 47.53 | 37.859 |
| P_VIGILAX | 106.931 | 131.265 | 197.611 |
| S_ATROMACULATUS | 1648.951 | 2005.182 | 2772.665 |
| S_VITREUS | 14.106 | 16.863 | 26.287 |

Appendix 11: Electronic data with PDF of maps and a shapefile which contain the results for all conservation planning results