

Figure 1: Life cycle diagram and census points for pre-reproduction census of gizzard shad.

Graphical Abstract

- An integral projection model for gizzard shad (Dorosoma cepedianum) utilizing
- density-dependent age-0 survival
- ⁴ J. Peirce^{1,4}, G. Sandland^{3,4}, B. Bennie¹, R.A. Erickson²

5

- 6 Highlights
- An integral projection model for gizzard shad (Dorosoma cepedianum) utilizing
- $_{8}$ density-dependent age-0 survival
- ⁹ J. Peirce^{1,4}, G. Sandland^{3,4}, B. Bennie¹, R.A. Erickson²
- Research highlight 1
- Research highlight 2

An integral projection model for gizzard shad (*Dorosoma cepedianum*) utilizing density-dependent age-0 survival

J. Peirce^{1,4}, G. Sandland^{3,4}, B. Bennie¹, R.A. Erickson²

1 University of Wisconsin - La Crosse, Mathematics & Statistics Department
2 U.S.G.S. Upper Mississippi Environmental Science Center
3 University of Wisconsin - La Crosse, Biology Department
4 River Studies Center

$_{19}$ Abstract

12

13

14

15

16

17

18

Gizzard shad (*Dorosoma cepedianum*) are a common freshwater fish found throughout the central and eastern portions of North America. Within these areas, gizzard shad play a number of critical roles in the freshwater community. Because of this, it is important that we understand how gizzard shad populations respond to environmental changes and what these changes may mean for aquatic communities in general and fish assemblages in particular. Here we introduce an integral projection model for gizzard shad based on empirical data and include density-dependent survival in age-0 fish. Integral projection models (IPM) are a generalization of stage-based, matrix population models that have been used to describe a wide range of organisms. IPMs are a natural choice for gizzard shad since many aspects of their life cycle have been studied. In this paper, we compared model outcomes to empirical patterns reported for this fish species at a key location along the Illinois River. Results of our work suggest that this model could serve as an important tool for predicting gizzard shad population responses to changing environmental conditions, including those mediated through species invasions.

- 20 Keywords: population dynamics, fisheries, Mississippi River basin, population ecology,
- 21 invasive species impact

2 1. Introduction

- Gizzard shad (*Dorosoma cepedianum*) is a laterally compressed, deep-bodied fish species
- that occupies numerous aquatic systems throughout central, southern and eastern regions
- of the United States (Pierce et al., 1981; Vanni et al., 2005). In more eutrophic habitats,

such as reservoirs, gizzard shad can reach high abunadances and can come to dominate fish assemblages. Because of this, gizzard shad have the potential to influence freshwater systems in a number of ways. First, young shad often serve as a critical food source for many fish species, including those of commercial and recreational importance (such as walleye and largemouth bass)(Jester et al., 1972). Thus, this species can serve as an important trophic link within aquatic food webs. Second, because detritus can serve as a primary food source throughout much of gizzard shad development (i.e. from the age-0 stage onward), these fish can transport nutrients from benthic regions into pelagic habitats (Mather et al., 1995; Schaus and Vanni, 2000; Vanni et al., 2005). This process can result in an increase in the nutrients available to organisms within the water column leading to increases in phytoplankton biomass, algal blooms, and, due to these conditions, shifts in freshwater community structure (Aday et al., 2003; Schaus and Vanni, 2000). Finally, the fact that detritus can comprise a substantial portion of gizzard shad diet also makes this species an important connection between terrestrial inputs and aquatic processes (Schaus and Vanni, 2000). Given its potentially important role in aquatic ecosystems, interest has intensified in understanding how gizzard shad populations respond to environmental changes (both natural and anthropogenic) and what these changes may mean for freshwater communities in general and fish assemblages in particular.

Interactions within and between species in interconnected environments can have important consequences for fish populations across space and time (Thorp et al., 2006). For
gizzard shad, previous work has suggested that fish densities can play an important role in
both the growth and survival patterns observed in these fish populations. For example, (Buynak et al., 1992) reported an inverse relationship between densities and the lengths of age-0
gizzard shad. Similarly (Welker et al., 1994) found that high densities of age-0 shad were
negatively associated with both fish length and survival under both field and semi-natural
conditions. Finally, (Michaletz, 2010) revealed that the densities of age-0 gizzard shad were
negatively correlated with survival in two Missouri lakes. These patterns were attributed
to intraspecific competition among young shad for prey (zooplankton) resources. Although
intraspecific competition may be influencing life-history traits in subsequent stages of gizzard
shad development, little work has actually been conducted to address this (DiCenzo et al.,

1996). There is also evidence that the densities of other co-occurring fish species (such as invasive carps) may also negatively influence aspects of gizzard shad biology, such as body condition (Irons et al., 2007; Love et al., 2018).

Although substantial empirical work on gizzard shad biology has accumulated over the decades, few if any studies have attempted to use these data to model the population dynamics of this species. Work by Catalano and Allen (2010, 2011) used empirically-based simulations of gizzard shad and focused on population-level responses. The authors did investigate population-structure using fish lengths but did not explore the effects of shad densities on population patterns. Here, we introduce an integral projection model for gizzard shad based on empirical data with density-dependent survival in age-0 fish. We then compare model outcomes to the dynamics reported for this fish species in the La Grange Station along the Illinois River. The model itself could be an important tool for predicting gizzard shad population responses to changing environmental conditions, including those mediated through species invasions (i.e., silver and bighead carp).

70 2. Model development

71 2.1. Gizzard shad life history

Mature gizzard shad tend to mate between May and June, although this can vary based on water temperatures. Males and females aggregate and then broadcast gametes into the surrounding water; fertilized eggs then settle and adhere to the bottom substrates. After a period of days, eggs hatch and fish develop from the larval stage to juveniles and eventually to adults. In many habitats, individuals can reach sexual maturity within a year. As gizzard shad mature, their diet preferences typically shift from phytoplankton and zooplankton early in development to detritus and zooplankton as adults. Given the large number of eggs produced by shad females (> 300,000/year), there is evidence that intraspecific competition can be intense during early developmental stages in this species. The strength of this competition may then subside as fish transition to feeding on different food items during latter stages of development.

Table 1: A summary of parameters, their biological meaning, and source for mean values.

| Parameter | Meaning (units) | Mean | Source |
|---------------------------------------|----------------------------------|---------------------------------------|----------------------------------|
| Logistic survival probability func- | | | |
| tion, $s(z)$ | | | |
| $s_{ m min}$ | minimum survival | 0.002 | (Bodola, 1955) |
| $s_{ m max}$ | maximum survival | $1 - 8.871K^{0.73}L_{\infty}^{-0.33}$ | THEN 2015 |
| α_s | inflection point | 104.34 | Estimated from LTRM dataset |
| eta_s | slope | -637.93 | Estimated from LTRM dataset |
| Growth function, $G(z, z')$ | | | |
| L_{∞} | maximum length (in mm) | 394.30 | (Catalano and Allen, 2010) |
| K_g | growth rate | 0.26 | (Michaletz, 2017) |
| σ_g | growth standard deviation | 25 | (Michaletz, 2017) |
| Normal distribution of length of age- | | | |
| $1, C_1(z')$ | | | |
| μ_r | mean length of recruitment (in | 105 | (Michaletz, 2017) |
| | mm) | | |
| σ_r | standard deviation of length | 25 | (Michaletz, 2017) |
| Eggs produced, $egg(z)$ | | | |
| $\mathrm{egg}_{\mathrm{max}}$ | maximum number of eggs pro- | 742,094 | Estimated from (Jons and Mi- |
| | duced | | randa, 1997) |
| α_e | inflection point | 314.44 | Estimated from (Jons and Mi- |
| | | | randa, 1997) |
| eta_e | slope | -7.12 | Estimated from (Jons and Mi- |
| | | | randa, 1997) |
| Survival of age-0, $s_0(d(t))$ | | | |
| a_0 | intercept | 0.27 | Estimated from (Michaletz, 2010) |
| b_0 | decay rate | 0.003 | Estimated from (Michaletz, 2010) |
| Spawning | | | |
| ν | probability that egg becomes vi- | 0.002 | (Bodola, 1955) |
| | able | | |
| p_b | probability that female spawns | 0.90 | |

83 2.2. Equations

We used an integral projection model to describe the life history of gizzard shad in the 84 Upper Mississippi River (UMR) system. Integral projection models (IPM) were introduced 85 by Easterling (Easterling et al., 2000), as a generalization of stage-based, matrix population models. IPMs have been used to describe a wide range of organisms (Ellner et al., 2016; Merow et al., 2014; Rees et al., 2014) and have only recently be used to model fish populations 88 (Erickson et al., 2017; Liao et al., 2019; White et al., 2016; Pollesch et al., 2022). The 89 availability of empirical size observation to aid with model parameterization makes IPM 90 a natural choice for gizzard shad since many aspects of their life cycle have been studied. 91 Specifically, functions used in our model incorporate data from studies on egg production and adult size, survival of the age-0 stage and density, and nearly thirty years of length 93 measurements in the main channel of the Upper Mississippi River system. 94

We assumed that variations among individual gizzard shad can be summarized by its length z (in mm) ranging from the minimum possible length L to the maximum value U. The state of the population at time t (in years) is described by the length distribution n(z,t). Specifically, for each time t, n(z,t) is a smooth function of z such that the number of individuals of length z in the interval [a,b] at time t is $\int_a^b n(z,t) \, dz$.

Between times t and t+1, individual gizzard shad may grow, die, and produce offspring 100 that vary in length depending on the individuals current length (Figure ??). At time t+1 the 101 population will have a length distribution defined by n(z, t+1). For our model, we partition 102 the life cycle of gizzard shad into two stages: 1) survival and growth, and 2) reproduction. 103 For an individual of length z at time t, $P(z',z)\Delta z$ is the probability that the individual 104 is alive at time t+1, and its size is in the interval $[z',z'+\Delta z]$ (as with n(z,t) this is an 105 approximation that is valid for small Δz , and the exact probability is given by an integral 106 like the one above). Similarly, $F(z',z)\Delta z$ is the number of new offspring in the interval 107 $[z', z' + \Delta z]$ present at time t + 1, per length-z individual at time t. 108

2.2.1. Growth and survival

109

We define P(z'z) = s(z,T)G(z',z) where s(z) is the adult annual survival probability and G(z',z) describes the annual length transitions. We assumed that the survival function

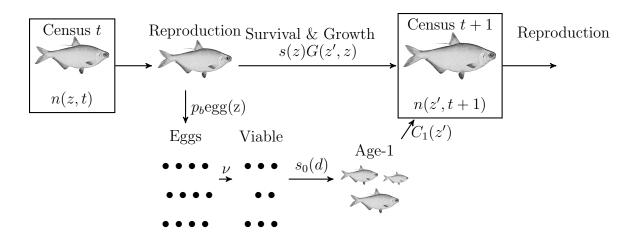


Figure 2: Life cycle diagram and census points for pre-reproduction census of gizzard shad.

is a logistic function,

$$s(z,T) = s_{\min} + \frac{s_{\max} - s_{\min}}{1 + e^{\beta_s(\ln(z) - \ln(\alpha_s))}}.$$
 (1)

with four parameters: the minimum survival rate s_{\min} ; a maximum survival rate, s_{\max} ; and intercept parameter, α_s ; and a slope parameter, β_s (Bolker, 2008).

We assumed that the growth function is a two-variable normal distribution centered around a modified von Bertalanffy function of the length at time t. The von Bertalanffy equation, commonly used to describe the length of a fish over time, is given by $z(t) = L_{\infty} \left(1 - e^{-K(t-t_0)}\right)$ where L_{∞} is maximum asymptotic length, K is the growth rate, and t_0 is the initial time. The expected length in the next year

$$z' = z(t+1) = L_{\infty} \left(1 - e^{-K(t+1-t_0)} \right) = L_{\infty} - L_{\infty} e^{-K(t-t_0)} e^{-K}$$
$$= L_{\infty} - (z(t) - L_{\infty}) e^{-K} = L_{\infty} \left(1 - e^{-K} \right) + z(t) e^{-K}.$$

Consequently, we assumed that $G(z',z) = \text{Prob}(z' | z, L_{\infty}, K_g) = \text{NormalPDF}(\mu_g, \sigma_g)$ where K_g is the individual growth rate, $\mu_g = L_{\infty} \left(1 - e^{-K_g}\right) + z(t)e^{-K_g}$, and σ_g is the standard deviation.

118 2.3. Fecundity

We define the fecundity kernel,

$$F(z',z) = p_b \exp(z) \nu s_0(n(z,t)) C_1(z')$$
(2)

where p_b is the probability of reproducing, egg(z) is the mean number of eggs produced, ν is the probability that an egg is viable, $s_0(n(z,t))$ is the density-dependent probability of surviving to age-1, and $C_1(z')$ is the length distribution of new recruits at age-1 (when they are first censused).

We assumed that the mean number of eggs produced by females of a certain length is a three-parameter logistic function,

$$\operatorname{egg}(z) = \frac{\operatorname{egg}_{\max}}{1 + e^{\beta_e(\ln(z) - \ln(\alpha_e))}}.$$
(3)

The probability of survival of gizzard shad during their first year may depend on many factors (Michaletz, 2010) such as mean temperature, mean total length, the present density of age-0 fish. In this study, we focus only on the density factor and define the probability of survival of age-0 fish as the exponential function

$$s_0(d(t)) = a_0 e^{-b_0 d(t)} (4)$$

where a_0 is the intercept, b_0 the decay rate, and d(t) is the density at time t of age-0 gizzard shad per 1000 m³,

$$d(t) = 10^{-3} \int_{L}^{U} p_b \operatorname{egg}(z) \nu n(z, t) dz.$$

Finally, after computing the total number of viable eggs produced and survive to age-1 fish, we multiplied this number with a normal distribution of length, $C_1(z') = \text{NormalPDF}(\mu_r, \sigma_r)$ where μ_r is the mean length of age-1 gizzard shad and σ_r is the standard deviation.

135 2.4. Dynamical model

The population at time t+1 is the sum of the contributions from each individual alive at time t,

$$n(z', t+1) = \int_{L}^{U} K(z', z) n(z, t) dz, \tag{5}$$

where K(z',z)=s(z)G(z',z)+F(z',z) and [L,U] is the range of all possible lengths.

3. Methods

3.1. Research area used for model parameterization

The Long Term Resource Monitoring (LTRM) is a U.S. Army Corp of Engineers program that assess, and detects changes in the fundamental health and condition of the Upper

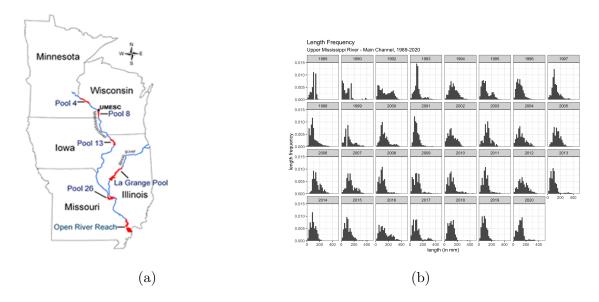


Figure 3: (a) Site source of image. (b) Length frequency of sampled gizzard shad in the main channel of the Upper Mississippi River system.

Mississippi River System (UMRS) ecosystem. Key ecological components of aquatic vegeta-143 tion, bathymetry, fish, land use/land cover, and water quality are monitored in the natural 144 floodplain between the head of navigation at Minneapolis, Minnesota and the confluence 145 with the Ohio River at Cairo, Illinois. The LTRM fish collection methodology includes a 146 multiple gear approach (netting and electrofishing) to monitor the general fish community in 147 six study pools/reaches through time (Gutreuter et al., 1995). Methodology, protocols and 148 modifications to the LTRM can be found in Gutreuter et al. (1995), and Ickes and Burkhardt 149 (2002). With the exception of La Grange Reach, five (Pools 4, 8, 13, 26, and the Open River 150 Reach) of the six study sites correspond to field stations located along the main channel of 151 the river (Figure 3a). Total length of gizzard shad in the main channel of the UMRS has 152 been recorded since 1989 with approximately 3000 random collections each year (Figure 3b). 153

3.2. Research area used for model validation

154

The 129 km long La Grange Reach is located between La Grange Lock and Dam (L&D) and Peoria L&D on the Illinois River, U.S., and is approximately midway between the Mississippi River and Lake Michigan. The Illinois River is a major tributary of the Mississippi River, draining nearly two-thirds of the state of Illinois. Along with the main channel of the UMRS, the fish community of La Grange Reach has been monitored by LTRM from 1990

to the present, with approximately 500 random collections each year from 15 June to 31 October.

The location of La Grange Reach has two important features motivation its choice for 162 the methods of our study. First, we were able to parameterized the IPM using data collected 163 from the main channel of the UMRS. As a part of Illinois River and UMRS, La Grange Reach is upstream and relatively independence from main channel of the Mississippi River. 165 In recent years, there have been concerns with the threatening introduction of invasive 166 carp to the Great Lakes. As a consequence, the impact of invasive carp on the native 167 fish populations in the pools leading to the Great Lakes have received an elevated level of 168 attention. Understanding the population dynamics of gizzard shad in La Grange Reach, may make it easier, in the future, to assess the impacts carp has on native fish populations. 170

3.3. Parameterization

3.3.1. Fecundity and recruitment

The maturity of female gizzard shad begin at approximately 140 mm and the number of 173 eggs that are produced increases as the females increase in size (Jons and Miranda, 1997). The logistic parameters for the mean number of eggs produced by females of a certain length 175 were obtained by fitting the three-parameter logistic function (Equation 3) to the data for 176 batch fecundity versus length (Jons and Miranda, 1997) (Figure 4a). The survival to age-0 177 was assumed to be dependent on the density of age-0 gizzard shad (Figure 4b). Parameters 178 for the exponentially decaying age-0 survival function (Equation 4) were determined by fitting equation 4 to the survival means for 2003-2007 cohorts of gizzard shard in five Missouri 180 reservoirs (Michaletz, 2010). To complete the recruitment process we assign a length to the 181 recruited individuals by simulating a Gaussian random variable with mean μ_c and standard 182 deviation σ_c . These parameters for the size distribution of age-1 fish were gleaned from 183 a study of gizzard shad located in large impoundments (Michaletz, 2017) and the historic 184 1990-2020 LTRM dataset of the main channel of the UMRS. 185

3.3.2. Growth and survival of adults

The parameters for the growth function were chosen as the mean values published on a study of gizzard shad located in large impoundments (Michaletz, 2017). The survival rate of

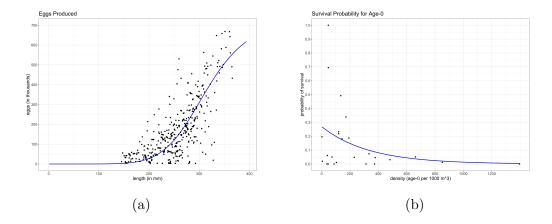


Figure 4: (a) Mean number of eggs produced by female gizzard shad egg(z). Data from (Jons and Miranda, 1997). (b) Age-0 density-dependent survival function $s_0(d)$. Data from (Michaletz, 2010).

adults gizzard shad dependent on their length is not well documented and required us to make 189 some additional modeling assumptions. We assume that the probability of adult survival is 190 related to the length by a four-parameter logistic function (Equation 1). An investigation of 191 gizzard shad in Lake Eerie (Bodola (1955)) provided the minimum and maximum survival 192 rate of adults. Based on the observed length distributions of gizzard shad sampled from the 193 main channel of the Mississippi River (Figure 3b), we assumed that solutions of our model 194 will exhibit periodic behavior every 8-9 years. We used a least squares method to estimate 195 the α_s and β_s parameters that minimized the total square-distance between the (observed) 196 pre-carp LTRM length distribution in the main channel of the UMRS and (predicted) model 197 equilibrium, n(z,t) during a 8-year period 100 years after initialization. The slope parameter 198 β_s was found to be large in magnitude resulting in a primarily two-valued survival probability. 199 Gizzard shad less than α_s mm in length have a very low survival rate (s_{\min}) while lengths 200 larger than α_s mm have the maximum survival rate s_{max} . This survival pattern has been 201 reported for a number of fish species and can arise for a number of reasons including feeding 202 attributes in predators (i.e. gape limitations, reduced abilities to catch prey, etc.) (Pepin et 203 al., 1992; Nowlin et al., 2006). 204

205 4. Analysis and results

We numerically solved the integral model using the Midpoint Rule with large approximating matrices (Burden and Faires, 2005). The Midpoint Rule has been commonly used

for integral projection models because of its simplicity and effectiveness (Ellner and Rees, 2006; Ramula et al., 2009; Merow et al., 2014). During the course of model development, we explored different step sizes for the Midpoint Rule and found that about 50 points provided numerically stable results. We integrated over lengths from 0 mm to 500 mm. The upper limit was chosen based upon numerical stability and consistency of the system (e.g., avoiding eviction or the loss of individuals due to numerical errors (Williams et al., 2012)).

214 4.1. Initial conditions

We assumed that the initial density of gizzard shad was $d_0 = 964.7$, the annual average density of gizzard shad observed in La Grange Reach from 1993-2019. The probability of an individual being length z at time t = 0 was assumed to be normally distributed with mean $0.5L_{\infty}$ and standard deviation $\sigma_0 = 30$. As a result, we initialized our model with length distribution

$$n(z,0) = d_0 \text{Norm}(0.5L_\infty, \sigma_0) = 964.7 \text{Norm}(166, 30).$$
 (6)

The model was coded in R (R Core Team, 2017) and the scripts are published on JP github page https://github.com/jppeirce.

222 4.1.1. Survival of age-0 cohort

The dependence on survival of next generation of age-0 fish on the present density of 223 age-0 fish strongly influences the density, at all ages, of gizzard shad within the population. 224 When the fish density is large, there may be more fish at longer lengths and consequently 225 a greater number of eggs produced. More eggs leads to a higher density of age-0 fish and reflectively a reduction in the survival to age-1. If this reduced survival continues for a 227 few years, the overall density of fish may decline and there may not be as many larger fish 228 reproducing. If fewer eggs are spawned, there are less age-0 fish and the reduced density 229 leads to a better survival probability. For these years we would expect the overall density to increase. This cycle of oscillation continues, reflected in our model by the time-dependent survival probability of age-0 recruits (Figure 5).

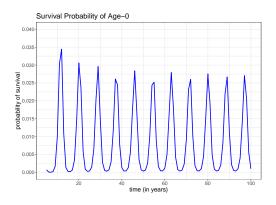


Figure 5

Figure 6: Survival probability of age-0 gizzard shad.

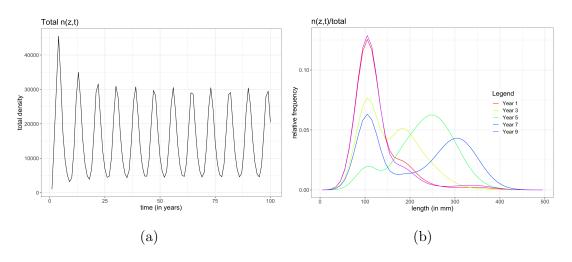


Figure 7: (a) The total number of gizzard shad in La Grange Reach predicted by IPM for first 100 years. (b) Simulated length distributions during an 8 year interval of time (approximately 1 period of the total density function)

233 4.1.2. Periodic orbit and validation with external dataset

234

235

236

237

238

The total number of gizzard shad in our simulation reached a stable periodic orbit (Figure 7a) within 50 years. The length distributions during the periodic orbit (Figure 7b) has similarities to length observations from La Grange found in the LTRM fish dataset (Figure 8a). In addition, the average simulated length distribution compares favorably to the length frequencies of observations from La Grange. (Figure 8b).

After simulating an additional 50 years, we fit a periodic function to the annual density of gizzard shad and determined a period of approximately 8.5 years. Figure 7b illustrates

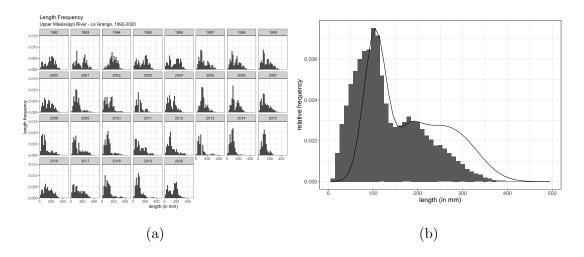


Figure 8: (a) Length frequencies of sampled gizzard shad in La Grange for each year, 1992-2020. (b) Length frequencies of sampled gizzard shad in La Grange from 1992-2020 compared with average (over 1 period) simulated length frequency.

the periodic length dynamics within the gizzard shad population during a 8-year window of the periodic orbit. The ebb and flow of the frequency of the age-1 cohort are associated 242 with the density-dependent survival function for age-0 fish as explained in Section 4.1.1. In 243 addition, Figure 7b captures the cohort length dynamics of the gizzard shad. Specifically, 244 the peak length frequency of age-1 fish in Year 1 near z = 105 mm is seen as a peak in Year 245 3 near 185 mm, a peak near 305 mm in Year 3, before becoming nearly unidentifiable in the graphs of Year 7 and 9. This is consistent with the lifespan reported for gizzard that 247 can range, on average, between 4 and 6 years (CITATION). The change in the location of 248 these relative maximum influences the number of eggs being produced and consequently the 249 density of age-0 fish the next year. An increase in the frequency of larger adults leads to 250 a delayed-reduction in the age-1 cohort as a result of our assumption of density-dependent 251 survival function. 252

5. Discussion

254

255

256

257

Gizzard shad are common in freshwater systems of North America where they are important components fo the freshwater community. Given this, it is important to understand how populations of this species vary with changing environmental circumstances, such as the occurrence of invasive species. Herein we use both LTRM data and parameters gleaned

from other empirical studies to develop an integral projection model for gizzard shad and test model outputs against patterns reported from a well-studied population of this species 259 from the Illinois River. 260

After fitting two adult survival parameters using LTRM data from the main channel 261 of the Mississippi River, we compared a simulated length distribution with LTRM data 262 collected from the La Grange Reach of the Illinois River from 1992 to 2020. The resulting 263 simulated length distributions (Figure 7b) reflected a number of the patterns observed in 264 gizzard shad captured from La Grange over the collection period (Figure 8a). Specifically, 265 the two peaks in Year 1 are similar to the observations made in 1997. In 1997, there is wider 266 range of mid-length and longer length gizzard shad and relative smaller amount of age-1 fish recorded. In the next year, there was a higher frequency of age-1 fish in both the LTRM 268 data and the simulations (Year 2). 269

We notice that the peak frequencies are near the same length with the model predicting 270 slightly more adults lengths and fewer juvenile lengths than the observations. This may be explained by gear bias. The methods used to capture fish make it more likely to record longer lengths (> 200mm). Studies [CITATION] suggest that due to the environmental 273 stochasticity and other effects, smaller recruitment fish densities can fluctuate annually and 274 be difficult to measure accurately. 275

271

While our model uses age-0 density to effect age-0 survival, we assumed constant viability 276 which may be sensitive to the external factors mentioned above. In addition, the location 277 of the maximum length and the variation in the of length of new recruits recorded in the 278 LTRM data, suggests that there may be smaller age-0 fish in La Grange Reach than in study 279 location (Michaletz, 2017) used to parameterize the model. 280

While not all observed length distributions are reflected in the simulations, the similarity 281 in the average length distributions (Figure 8b) suggest that our IPM could serve as a tool for predicting length distributions for gizzard shad across different study areas. (LAST 283 SENTENCE NEEDS A LOT MORE CLARITY) 284

Gaining an understanding of how length distributions of gizzard shad emerge under 285 density-dependent survival in the age-0 class will serve as a foundation for investigating 286 density effects at subsequent stages in the life cycle. In addition, this single-species model could also be expanded to incorporate interspecific interactions between gizzard shad and species such as invasive carp, which appear to negatively impact gizzard shad life-histories through competition for food resources.

291 6. Acknowledgments

These data are a product of the U.S. Army Corps of Engineer's Upper Mississippi River 292 Restoration Program (UMRR) Long Term Resource Monitoring (LTRM) element imple-293 mented by the U.S. Geological Survey in collaboration with the five Upper Mississippi River 294 System (UMRS) states of Illinois, Iowa, Minnesota, Missouri, and Wisconsin. The U.S. 295 Army Corps of Engineers (Corps) provides guidance and has overall program responsibility. 296 We thank the U.S. Geological Survey Biothreats program and Great Lakes Restoration 297 Initiative for funding. In addition, research was supported by NSF-DMS Grant #1852224, 298 "REU Site: Ecological Modeling of the Mississippi River Basin". 299

300 References

- R. J. Pierce, T. E. Wissing, B. A. Megrey, Aspects of the feeding ecology of gizzard shad in

 Acton Lake, Ohio, Transactions of the American Fisheries Society 110 (3) (1981) 391–395.
- M. J. Vanni, K. K. Arend, M. T. Bremigan, D. B. Bunnell, J. E. Garvey, M. J. Gonzalez,
 W. H. Renwick, P. A. Soranno, R. A. Stein, Linking landscapes and food webs: effects of
 omnivorous fish and watersheds on reservoir ecosystems, BioScience 55 (2) (2005) 155–167.
- D. B. Jester, B. L. Jensen, et al., Life history and ecology of the gizzard shad, *Dorosoma* cepedianum (Le Sueur) with reference to Elephant Butte Lake .
- M. E. Mather, M. J. Vanni, T. E. Wissing, S. A. Davis, M. H. Schaus, Regeneration of nitrogen and phosphorus by bluegill and gizzard shad: effect of feeding history, Canadian Journal of Fisheries and Aquatic Sciences 52 (11) (1995) 2327–2338.
- M. H. Schaus, M. J. Vanni, Effects of gizzard shad on phytoplankton and nutrient dynamics: role of sediment feeding and fish size, Ecology 81 (6) (2000) 1701–1719.

- D. D. Aday, R. J. H. Hoxmeier, D. H. Wahl, Direct and indirect effects of gizzard shad on bluegill growth and population size structure, Transactions of the American Fisheries Society 132 (1) (2003) 47–56.
- J. H. Thorp, M. C. Thoms, M. D. Delong, The riverine ecosystem synthesis: biocomplexity in river networks across space and time, River Research and Applications 22 (2) (2006) 123–147.
- G. L. Buynak, R. S. Hale, B. Mitchell, Differential growth of young-of-year gizzard shad in several Kentucky reservoirs, North American Journal of Fisheries Management 12 (3) (1992) 656–662.
- M. T. Welker, C. L. Pierce, D. H. Wahl, Growth and survival of larval fishes: roles of competition and zooplankton abundance, Transactions of the American Fisheries Society 123 (5) (1994) 703–717.
- V. J. DiCenzo, M. J. Maceina, M. R. Stimpert, Relations between reservoir trophic state and gizzard shad population characteristics in Alabama reservoirs, North American journal of fisheries management 16 (4) (1996) 888–895.
- K. S. Irons, G. Sass, M. McClelland, J. Stafford, Reduced condition factor of two native fish species coincident with invasion of non-native Asian carps in the Illinois River, USA Is this evidence for competition and reduced fitness?, Journal of Fish Biology 71 (2007) 258–273.
- S. A. Love, N. J. Lederman, R. L. Anderson, J. A. DeBoer, A. F. Casper, Does aquatic invasive species removal benefit native fish? The response of gizzard shad (Dorosoma cepedianum) to commercial harvest of bighead carp (Hypophthalmichthys nobilis) and silver carp (H. molitrix), Hydrobiologia 817 (1) (2018) 403–412.
- M. J. Catalano, M. S. Allen, A size-and age-structured model to estimate fish recruitment, growth, mortality, and gear selectivity, Fisheries Research 105 (1) (2010) 38–45.
- M. J. Catalano, M. S. Allen, A whole-lake density reduction to assess compensatory responses

- of gizzard shad Dorosoma cepedianum, Canadian Journal of Fisheries and Aquatic Sciences
 68 (6) (2011) 955–968.
- A. Bodola, The life history of the gizzard shad, Dorosoma cepedianum, in western Lake Erie,
 Ph.D. thesis, Doctoral dissertation. Ohio State University, Columbus, 1955.
- P. Michaletz, Variation in characteristics among gizzard shad populations: The role of impoundment size and productivity, Fisheries Management and Ecology 24 (5) (2017) 361–371.
- G. Jons, L. Miranda, Ovarian weight as an index of fecundity, maturity, and spawning periodicity, Journal of Fish Biology 50 (1) (1997) 150–156.
- P. H. Michaletz, Overwinter survival of age-0 gizzard shad in Missouri reservoirs spanning a productivity gradient: roles of body size and winter severity, Transactions of the American Fisheries Society 139 (1) (2010) 241–256.
- M. R. Easterling, S. P. Ellner, P. M. Dixon, Size-specific sensitivity: applying a new structured population model, Ecology 81 (3) (2000) 694–708.
- S. P. Ellner, D. Z. Childs, M. Rees, et al., Data-driven modelling of structured populations,
 A practical guide to the Integral Projection Model. Cham: Springer.
- C. Merow, J. P. Dahlgren, C. J. E. Metcalf, D. Z. Childs, M. E. Evans, E. Jongejans,
 S. Record, M. Rees, R. Salguero-Gómez, S. M. McMahon, Advancing population ecology
 with integral projection models: a practical guide, Methods in Ecology and Evolution
 5 (2) (2014) 99–110.
- M. Rees, D. Z. Childs, S. P. Ellner, Building integral projection models: a user's guide, Journal of Animal Ecology 83 (3) (2014) 528–545.
- R. A. Erickson, E. A. Eager, M. K. Brey, M. J. Hansen, P. M. Kocovsky, An integral projection model with YY-males and application to evaluating grass carp control, Ecological Modelling 361 (2017) 14–25.

- B. Liao, X. Shan, C. Zhou, Y. Han, Y. Chen, Q. Liu, A dynamic energy budget-integral
- projection model (DEB-IPM) to predict population-level dynamics based on individual
- data: a case study using the small and rapidly reproducing species Engraulis japonicus,
- Marine and Freshwater Research 71 (4) (2019) 461–468.
- J. W. White, K. J. Nickols, D. Malone, M. H. Carr, R. M. Starr, F. Cordoleani, M. L.
- Baskett, A. Hastings, L. W. Botsford, Fitting state-space integral projection models to
- size-structured time series data to estimate unknown parameters, Ecological Applications
- 26 (8) (2016) 2677–2694.
- N. Pollesch, K. Flynn, S. Kadlec, J. Swintek, S. Raimondo, M. Etterson, Developing integral
- projection models for ecotoxicology, Ecological Modelling 464 (2022) 109813.
- B. M. Bolker, Ecological models and data in R, Princeton University Press, 2008.
- S. Gutreuter, R. Burkhardt, K. S. Lubinski, Long Term Resource Monitoring Program procedures [microform] .
- B. S. Ickes, R. W. Burkhardt, Evaluation and proposed refinement of the sampling design
- for the Long Term Resource Monitoring Program's fish component, Tech. Rep., Geological
- Survey La Crosse WI Upper Midwest Environmental Sciences Center, 2002.
- R. Burden, J. D. Faires, Numerical analysis 8th ed, Thomson Brooks/Cole.
- S. P. Ellner, M. Rees, Integral projection models for species with complex demography, The
- 382 American Naturalist 167 (3) (2006) 410–428.
- 383 S. Ramula, M. Rees, Y. M. Buckley, Integral projection models perform better for small
- demographic data sets than matrix population models: a case study of two perennial
- herbs, Journal of Applied Ecology 46 (5) (2009) 1048–1053.
- J. L. Williams, T. E. Miller, S. P. Ellner, Avoiding unintentional eviction from integral
- projection models, Ecology 93 (9) (2012) 2008–2014.
- R Core Team, R: A Language and Environment for Statistical Computing, R Foundation
- for Statistical Computing, Vienna, Austria, URL https://www.R-project.org/, 2017.