Applying a reservoir functional-zone paradigm to littoral bluegills: differences in length and catch frequency?

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ABSTRACT

Reservoirs exhibit gradients in conditions and resources along the transition from lotic to lentic habitat that may be important to bluegill ecology. The lotic–lentic gradient can be partitioned into three functional zones: the riverine, transitional, and lacustrine zones. We measured catch frequency and length of bluegills (*Lepomis macrochirus*) captured along the periphery of these areas (i.e., in the littoral zone of each functional zone) for four small reservoirs in Southeastern Ohio during the summer months of three years. Catch frequency differed between zones for two reservoirs, but these differences were not observed in other years. There was no relationship between reservoir zone and either standard length or catch frequency when the data for all reservoirs were pooled, but we did observe a bimodal length distribution in all reservoirs. A combination of ecological factors including inter and intraspecific competition, predation intensity, management practices, limnology, and assemblage complexity may be mitigating bluegill distribution and abundance in reservoirs. Therefore, a functional zone (categorical) approach to understanding bluegill ecology in reservoirs may not be appropriate.

INTRODUCTION

Bluegill (*Lepomis macrochirus*) exhibit ontogenetic habitat shifts that coincide with shifts in foraging behavior in natural lakes. After hatching in the littoral zone, young-of-year bluegill migrate to the limnetic zone to feed on zooplankton (*Werner, 1969*). Once a larger body size has been attained, bluegill return to the littoral zone and feed opportunistically amongst macrophytes. After several years feeding in the littoral zone, larger bluegills shift back to a diet of zooplankton and move freely between the littoral and limnetic zones (*Mittelbach, 1981*). Shifts in bluegill diet and habitat use may be a result of a trade-off between maximizing foraging efficiency while minimizing predation risk (*Werner & Hall, 1988*). However, *Wildhaber & Lamberson (2004)* suggested an alternative hypothesis based on a hierarchical model of tradeoffs between prey availability and temperature in lakes. Regardless of the specific cause of the shift in bluegill habitat use (direct selection pressure...
via predation vs. indirect selection pressure from prey/habitat availability), it is an effective life history strategy (reviewed by Werner & Peacor, 2003).

The success of habitat switching as a life history strategy for bluegill may depend on a number of factors. For instance, basin morphometry may lead to differential recruitment success of bluegill amongst natural lakes where maximum depth, percent littoral area (Tomcko & Pierce, 2001), and lake surface area (Tomcko & Pierce, 2005) have all been linked to recruitment success. Littoral bluegill abundance is positively associated with habitat features such as the availability of woody debris (Newbrey et al., 2005), and native macrophytes (Theel & Dibble, 2008). Another important factor is the availability of zooplankton (Garvey & Stein, 1998); lakes with low productivity or high turbidity have low epilimnetic phytoplankton abundance which reduces zooplankton and thus decreases bluegill recruitment (Stein, DeVries & Dettmers, 1995). High abiotic turbidity in the photic zone is normally driven by physical processes such as wind mixing and flooding but also can be influenced by sympatric species that resuspend sediment (e.g., gizzard shad; Vanni et al., 2005) resulting in both direct and indirect density-dependent effects on bluegill recruitment via alteration in prey availability and/or capture success (Aday & Hoxmeier, 2003; Shoup et al., 2007). Indeed, protracted spawning by bluegill (Garvey, Herrera & Leggett, 2002; Kaemingk et al., 2014) may be an adaptation to offset density-dependent effects caused by competition for prey (Partridge & DeVries, 1999; Michaletz, 2006; but see Leonard, DeVries & Wright, 2010).

Reservoirs exhibit gradients in the relative area of littoral vs. limnetic habitat (Thornton, 1990), zooplankton community composition (Bernot et al., 2004), and a suite of environmental variables including turbidity (Thornton, 1990) and available nutrients (Kennedy & Walker, 1990) vary along the lotic-lentic transition. Reservoirs can be divided into three functional zones based on these gradients (Fig. 1): the fluvial zone is the shallow unstratified portion that is heavily influenced by flooding and where well-mixed epilimnetic water is in direct contact with sediments, the transitional zone is weakly stratified and less influenced by flooding or sediment resuspension, and the lacustrine zone is the stably stratified lake-like area (adapted from Kimmel, Lind & Paulson, 1990).

Interpreting the ecological dynamics of reservoirs in the paradigm of functional zones along the lotic-lentic transition has been regularly applied to organisms that are at the whim of hydrologic conditions (reviewed by Ruhl, 2013), but to our knowledge has not
Table 1  Basin morphometrics, fill date, and date of last bluegill stocking for each reservoir.

| Variable                  | Dow   | Fox   | Hope  | Snowden |
|---------------------------|-------|-------|-------|---------|
| Catchment area (km$^2$)   | 18.90 | 10.36 | 25.64 | 9.78    |
| Surface area (km$^2$)     | 0.67  | 0.23  | 0.48  | 0.65    |
| Maximum depth (m)         | 9.5   | 6.0   | 6.5   | 10.0    |
| Mean depth (m)            | 1.62  | 1.28  | 1.31  | 2.45    |
| Volume (m$^3$)            | 1,085,069 | 2,94,975 | 630,744 | 1,590,558 |
| Shoreline length (km)     | 11.27 | 3.86  | 9.18  | 11.91   |
| Shoreline development     | 5.49  | 3.21  | 5.29  | 5.89    |
| Maximum fetch (km)        | 2.00  | 0.71  | 1.07  | 2.87    |
| Fill date                 | 1960  | 1968  | 1939  | 1970    |
| Last stocked with bluegill| 1972  | N/A   | 1979  | 1970    |

been explicitly assessed in relation to more motile species such as fish. Additionally, the functional zonation paradigm for reservoirs typically describes open water rather than along the shoreline (littoral zone), despite the fact that differences in the mixing regime in open water may directly influence factors such as nutrient availability along the periphery. Because bluegill ecology is intimately linked to the conditions and resources in the limnetic as well as littoral zones, the functional zone paradigm may be particularly relevant and yield insight into broadscale differences in their ecology within reservoirs (i.e., both along the lotic-lentic gradient and between the littoral and limnetic zones). Specifically, we predicted that size and catch frequency may vary among functional zones because of differences that affect bluegill recruitment (i.e., their suitability for growth and reproduction, see above). We sampled bluegill abundance in the littoral zone throughout four different reservoirs to assess their use of the lotic-lentic transition, (in multiple years in some cases) during July and August when stable thermal stratification is normally strongest and therefore differences among functional zones may be at their peak.

METHODS

Study sites

We sampled Dow Lake, Lake Hope, Lake Snowden and Fox Lake. These four reservoirs are located in close proximity to one another in the unglaciated hills of Southeastern Ohio and are managed by units of the Ohio Department of Natural Resources (Fig. 2, Table 1).

Dow Lake (Stroud’s Run State Park) is used primarily for recreation, but also to mitigate flooding of the Hocking River downstream of Athens, Ohio. This reservoir was filled during 1960 and the watershed is composed of minimally disturbed hills, woodland, and open fields. Throughout the reservoir, the littoral zone has been modified via the felling of shoreline trees and addition of brush piles to coves during the early 2000s (M Greenlee, District 4 ODNR Biologist, pers. comm., 2010).

Lake Snowden was filled during 1970. The reservoir previously supplied drinking water to the surrounding community, but currently is used for flood control, hatchery water supply, and recreational activities. The watershed consists of rolling hills, agricultural
fields, and woodlots while the shoreline habitat includes submerged trees, overhanging brush and abundant submerged macrophytes.

Fox Lake was filled during 1968 and the watershed is composed of rolling hills, agricultural fields and woodlots. High sedimentation rates in the riverine zone have resulted in poor angler access to the reservoir and, consequently, submerged macrophytes were removed during the mid-1990’s in order to increase flow and accessibility to the riverine zone (M Greenlee, District 4 ODNR Biologist, pers. comm., 2010). These efforts have not been successful in improving angler access and dredging to manually remove sediment is impractical for this reservoir.

Lake Hope is located within the Zaleski State Forest and was filled during 1937. The watershed is composed of mature second growth forest scattered with abandoned pit and shaft coal mines. The reservoir has abundant invasive emergent macrophytes (primarily fragrant water lily, *Nymphaea odorata*) mixed with a variety of other emergent and submerged macrophytes around the periphery.

Bluegill are not regularly stocked into any of the reservoirs (Table 1). Rainbow trout (*Oncorhynchus mykiss*, mean length 303 mm, 2001–2011) are stocked into Dow Lake every
Table 2  Summary of the number of sampling sites and the total catch for each reservoir, zone, and year that sampling occurred.

| Reservoir/year | Zone        | #Sites | Total catch |
|----------------|-------------|--------|-------------|
| Dow 2006       | Riverine    | 4      | 57          |
|                | Transitional| 4      | 22          |
|                | Lacustrine  | 8      | 67          |
| Fox 2006       | Riverine    | 5      | 72          |
|                | Transitional| 2      | 19          |
|                | Lacustrine  | 2      | 27          |
| Dow 2007       | Riverine    | 17     | 104         |
|                | Transitional| 11     | 103         |
|                | Lacustrine  | 12     | 92          |
| Hope 2007      | Riverine    | 4      | 31          |
|                | Transitional| 5      | 47          |
|                | Lacustrine  | 6      | 60          |
| Snowden 2007   | Riverine    | 3      | 25          |
|                | Transitional| 6      | 59          |
|                | Lacustrine  | 7      | 40          |
| Hope 2008      | Riverine    | 4      | 19          |
|                | Transitional| 6      | 47          |
|                | Lacustrine  | 10     | 43          |

Notes.
Raw data were used for statistical analysis (not the means presented here).

April. All four reservoirs are stocked yearly or in alternating years with channel catfish (*Ictalurus punctatus*, mean 221 mm) during the Fall. Lakes Snowden and Hope are stocked with saugeye (*Sander canadensis × Sander vitreus*, mean 31.5 mm) every year during the Spring. Fish are normally stocked into the reservoirs in close proximity to the boat launch, meaning that stocked fish are introduced into the riverine zone at Dow Lake and Fox Lake, the transitional zone at Lake Snowden, and the lacustrine zone at Lake Hope.

Sampling regime
The reservoirs were sampled using shore-line traps during three different years, but only Dow Lake and Lake Hope were repeatedly sampled (Table 2). Trapping was conducted during July and August in all years, but the number of weeks during which trapping occurred varied by year. All trapping was conducted using a randomized block design both within and among reservoirs, thereby minimizing the likelihood of a temporal effect among reservoirs or reservoir zones within a given year. Sampling methods were in accordance with Ohio University IACUC protocols and Ohio Department of Natural Resources Permit #464.

At each trapping site, pairs of oval traps (Promar ‘large’ 81 × 50 × 30 cm (1 cm mesh size and 12 cm minimum tunnel diameter) and ‘extra-large’ 91 × 62 × 50 cm (2.5 cm mesh and 15 cm tunnel diameter)) were positioned approximately 2 m from one another with trap entrances positioned parallel with the shoreline. The distance that the traps were positioned from shore was dictated by the slope of the shoreline; in order to avoid
drowning concurrently caught turtles. The traps were positioned such that a small portion of trap protruded from the surface. Each site used two ‘large’ traps during the 2006 sampling season. During the 2007 and 2008 sampling season each site had one ‘large’ and one ‘extra-large’ trap. ‘Extra-large’ traps were introduced during 2007 and 2008 to ensure that we were not excluding larger bluegills and thus validate the 2006 size data. Trapping sites were located at approximately equal intervals around the periphery (littoral zone) of each reservoir. Upon arriving at the pre-determined trapping location, the exact positioning of the traps was again dictated by the slope of the shoreline. Each trap was baited with commercially available dip bait (Premo brand ‘original super-sticky dip bait’) hung inside the trap in a cheesecloth bag and positioned in as flat a position as the shoreline allowed in order to keep turtles and fish from getting under the trap or causing the trap to shift into deeper water. We checked traps every 24 h for five days and recorded species and standard length of all fish and then released them immediately.

Analysis
We determined the transitional zone area and the size of the riverine and lacustrine zones a posteriori for each reservoir and each year. We defined the transitional zone as the area of the reservoir where the presence of thermal stratification fluctuated due to weather conditions (wind and flooding) during the sampling period. Therefore, the transitional zone begins at the point when a well-mixed epilimnion and a metalimnion are present outside of the thalweg (if present) and continues until underflows terminate into interflows through the metalimnion (Fig. 1).

Length and catch frequency (the total number of fish caught during a five-day period for each site) could not be normalized for all groups, so comparisons among reservoir zones (i.e., within each reservoir) were conducted using Kruskal–Wallis tests and a priori Mann–Whitney U tests. The same tests were used when comparing catch frequency among reservoir zones for all reservoirs combined, but one-way ANOVA with post-hoc Tukey tests were employed to compare the standard length among zones for all reservoirs combined. Although the length data were not normal, ANOVA is robust for non-parametric data at sample sizes of 30 or greater for each group when the model is balanced. For our unbalanced model, normality can generally be assumed at sample sizes greater than 100 per group and our sample sizes are approximately 300 per group. We compared the catch frequency of small vs. large bluegills among reservoir zones for the pooled data (all reservoirs combined) using a two sample Kolmogorov–Smirnov test. All statistics were performed using SPSS 12.0.

RESULTS
Standard length
Standard length only varied by reservoir zone for Dow Lake during 2006. In that case, bluegill caught in the transitional zone were smaller than those caught in the other zones (riverine: Mann–Whitney, \( U = 412, p = 0.019 \); lacustrine: \( U = 431.5, p = 0.004 \)), but this result was not seen during 2007 (Fig. 3). When the length data from all reservoirs was
Figure 3 Box-plot of bluegill standard length between zones at Dow Lake during 2006 and 2007. Bluegills caught in the transitional zone during 2006 were significantly smaller than those caught in the riverine (Mann–Whitney, \( U = 412, p = 0.019 \)) or lacustrine (\( U = 431.500, p = 0.004 \)) zones, but this result was not observed during 2007. Box represents first and third quartiles, whiskers positioned at ±2 SD, horizontal line is the mean.

combined, there were no differences among zones (one-way ANOVA, \( F_{(2, 822)} = 0.053, p = 0.921 \)).

**Catch frequency**

There was no difference in the catch frequency of bluegill among reservoir zones for any of the reservoirs (Table 3). Catch frequency did not vary among reservoir zones when the data from all reservoirs was pooled either (Kruskal–Wallis, \( \chi^2 = 1.094, p = 0.579 \)). Because the distribution of lengths was bimodal for all reservoirs in all years, we split the dataset at the saddle of the distribution (>/>8.5 cm, Fig. 4) and asked if the number of small or large bluegills varied over reservoir zone for each reservoir. Only in Lake Hope were small bluegills encountered more often in the transitional zone than in the fluvial zone during 2008 (Mann–Whitney, \( U = 357, p = 0.019 \), Fig. 5), but this result was not observed during the previous year. Combined for all reservoirs, there was no difference
Figure 4 Histogram of bluegill standard lengths for all reservoirs combined. The dashed line indicates the saddle in the distribution at 8.5 cm where the data was bifurcated into “small” and “large”.

Table 3 Results of a Kruskal–Wallis analysis of catch frequency between reservoir zones for each reservoir and year.

| Lake     | Year | $\chi^2$ | p-value |
|----------|------|----------|---------|
| Fox      | 2006 | 2.881    | 0.237   |
| Dow      | 2006 | 5.094    | 0.078   |
| Dow      | 2007 | 0.550    | 0.760   |
| Snowden  | 2007 | 0.793    | 0.673   |
| Hope     | 2007 | 0.812    | 0.666   |
| Hope     | 2008 | 1.832    | 0.400   |

in the catch frequency of both small and large bluegills among zones (Kruskal–Wallis; small: $\chi^2 = 2.285, p = 0.319$; large: $\chi^2 = 0.406, p = 0.816$). Additionally, there was no relationship between the catch frequency of small vs. large bluegills among reservoir zones (Kolmogorov–Smirnov, $Z = 1.083, p = 0.192$).
DISCUSSION

Bluegill populations are influenced by a variety of factors including both abiotic factors such as turbidity (Stein, DeVries & Dettmers, 1995) or temperature (Wildhaber & Lamberson, 2004) and biotic factors such as prey availability (Garvey & Stein, 1998; Hoxmeier, Aday & Wahl, 2009) or predators (Werner & Hall, 1988); all of these factors vary dramatically among reservoir zones as a simple function of stratification regime (as well as other factors such as nutrient loading, water retention time, etc.) (Kimmel, Lind & Paulson, 1990). However, few differences in bluegill among reservoir zones were observed in our study. Size of bluegill differed among zones at Dow Lake during 2006, but this result was not observed during 2007. Similarly, small bluegills were caught more frequently in the transitional zone at Lake Hope during 2008, but not during 2007. When the data from all reservoirs was pooled, there were no differences in either size or catch frequency among reservoir zones, suggesting that habitat partitioning may be based on different criteria in reservoirs (Gelwick & Matthews, 1990; Eggleton et al., 2005) than has previously been described for natural lakes (e.g., Werner et al., 1977).

The lack of repeatability in our findings among years may be indicative of the true nature of reservoirs as a habitat for bluegill. Resources and conditions within a reservoir may be dependent on prevailing weather patterns (Lienesch & Matthews, 2000; but see Edwards et al., 2007), inputs from the watershed (Gido et al., 2002; Vanni et al., 2005) and
presence of certain species (e.g., gizzard shad; Vanni et al., 2005). All of these variables can fluctuate dramatically year to year and cause shifts in prey availability (Betsill & van den Avyle, 1994) and predation intensity (Jackson & Noble, 2000). Additionally, the artificial, managed nature of reservoirs creates dynamic environments where water levels and habitat availability/suitability (Collingsworth & Kohler, 2010) and stocking of competitors (Leonard, DeVries & Wright, 2010) and/or predators may vary yearly. Therefore, while size and catch frequency of bluegill may differ by reservoir zone at times (as we observed at Dow during 2006 and Hope during 2008), they are likely influenced by other factors as well, which may have disrupted our ability to consistently detect differences among zones.

Bluegill spawning behavior also may influence the detectability of differences in length and catch frequency among reservoir zones (if they exist). Bluegill spawning is condition-dependent for males (males in better physical condition spawn first; Cargnelli & Neff, 2006), which results in protracted spawning (spawning over an extended period; Kaemingk et al., 2014). Given the differences in prey availability among reservoir zones (Betsill & van den Avyle, 1994), protracted spawning may be more prevalent in reservoirs than in lakes and could cause behavioral plasticity in habitat use that is difficult to detect using standard techniques (e.g., trapping, netting, or electro-shocking). That is, if bluegill spawning occurs over a wider range of times in reservoirs, population-wide shifts in habitat use would be similarly spread over a longer duration and differences among zones, which may be important to bluegill, may also be difficult to detect. This is supported by Jolley, Edwards & Willis (2009) who found that the timing of spawning in bluegill varied among nearby reservoirs and among years in the same reservoirs.

The size structure of the bluegills we caught by trapping (all reservoirs combined) was bimodal and somewhat positively skewed. The positive skew was determined by the uniformly smaller bluegill that freely travel through the traps without being caught while the right tail extends because larger individuals are rare. The saddle in the size distribution at approximately 8.5 cm is more intriguing. Bluegills < 10 cm (except planktivorous larvae) are normally found in the littoral zone of lakes because this area provides the greatest protection from predation (Werner & Hall, 1979). It may be that in our study, bluegill move away from the shoreline reservoir-wide at a much smaller size in reservoirs than in natural lakes, but we believe this is unlikely given the differences in ‘offshore’ conditions and resources among reservoir zones. Likewise, it is possible that the saddle of the distribution represents two different age classes, but this is also unlikely given the variation in growth rates observed in bluegills among reservoirs (Jackson, Quist & Larscheid, 2008) and their protracted spawning behavior. More likely, the saddle is a result of size selective predation by largemouth bass (Olson, 1996) or other piscivores such as saugeye. Because these piscivores are gape limited, bluegills over approximately 10 cm (Werner & Hall, 1979) are at lower risk of predation than smaller bluegills (Santucci & Wahl, 2003). Therefore, the saddle may represent the point at which size-specific mortality of bluegill caused by predation (Mittelbach & Persson, 1998) starts to decline in Southeastern Ohio reservoirs.
Lastly, another factor that may have contributed to our results is that our methodology did not detect temporal variation within a reservoir. Because trapping occurred during the course of a few weeks for each reservoir, differences in catch frequency or size among zones as a result of behavioral plasticity during ontogeny may be diluted. However, Gelwick & Matthews (1990) suggest that there is little temporal variation in littoral fish assemblages of reservoirs relative to lakes because these assemblages are ‘evolutionarily short-lived’. That is, because a given reservoir has not existed long in evolutionary time, fish assemblages may not exhibit the same patterns seen in natural lakes which have existed for many years. Our results seem to support this conclusion given that we only saw differences in the oldest of the reservoirs we sampled. Similarly, anthropogenic factors such as intensive stocking (Gelwick & Matthews, 1990) or the maintenance of a community dominated by a small number of species (Eggleton et al., 2005) may contribute to a decrease in temporal variation in habitat use in reservoirs.

In this study, bluegill generally did not differ in size or catch frequency among reservoir zones in four Southeastern Ohio reservoirs. This result, although unexpected due to the broad differences in habitat characteristics among reservoir zones, may be caused by a combination of factors including prey availability relative to predation intensity in reservoirs, management practices, limnology, and assemblage complexity. Kaemingk et al. (2014), working with limnetic bluegills, hypothesized that similar factors may regulate the timing and duration of spawning behavior, which should have reinforced differences in bluegill abundance among reservoir zones. Despite the fact that significant differences were found using the same approach with aquatic turtles (Ruhl, 2013), we did not detect differences in bluegill size or catch frequency among zones. Thus, it is likely that a categorical (functional zone) approach to detecting differences in bluegill ecology within a reservoir is not appropriate.

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Competing Interests
The authors declare there are no competing interests.
Author Contributions

- Nathan Ruhl conceived and designed the experiments, performed the experiments, analyzed the data, contributed reagents/materials/analysis tools, wrote the paper, prepared figures and/or tables, reviewed drafts of the paper.
- Holly DeAngelis and Abigale M. Crosby analyzed the data, wrote the paper, prepared figures and/or tables, reviewed drafts of the paper.
- Willem M. Roosenburg analyzed the data, wrote the paper, reviewed drafts of the paper.

Animal Ethics

The following information was supplied relating to ethical approvals (i.e., approving body and any reference numbers):

This study was conducted in accordance with Ohio University’s IACUC guidelines.

Field Study Permissions

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REFERENCES

Aday DD, Hoxmeier JH, Wahl DH. 2003. Direct and indirect effects of gizzard shad on bluegill growth and population size structure. Transactions of the American Fisheries Society 132:47–56 DOI 10.1577/1548-8659(2003)132<0047:DAIEOG>2.0.CO;2.

Bernot RJ, Dodds WK, Quist MC, Guy CS. 2004. Spatial and temporal variability of zooplankton in a great plains reservoir. Hydrobiologia 525(1–3):101–112 DOI 10.1023/B:HYDR.0000038857.19342.fd.

Betsill RK, van den Avyle MJ. 1994. Spatial heterogeneity of reservoir zooplankton—a matter of timing. Hydrobiologia 277(1):63–70 DOI 10.1007/BF00023986.

Cargnelli LM, Neff BD. 2006. Condition-dependent nesting in bluegill sunfish Lepomis macrochirus. Journal of Animal Ecology 75(3):627–633 DOI 10.1111/j.1365-2656.2006.01083.x.

Collingsworth PD, Kohler CC. 2010. Abundance and habitat use of juvenile sunfish among different macrophyte stands. Lake and Reservoir Management 26(1):35–42 DOI 10.1080/07370651003634380.

Edwards KR, Phelps QE, Graeb BDS, Willis DW. 2007. Asynchronous bluegill recruitment in four South Dakota impoundments. Journal of Freshwater Ecology 22(1):19–22 DOI 10.1080/02705060.2007.9664141.

Eggleton MA, Ramirez R, Hargrace CW, Gido KB, Masoner JR, Schnell GD, Matthews WJ. 2005. Predictability of littoral-zone fish communities through ontogeny in Lake Texoma, Oklahoma-Texas, USA. Environmental Biology of Fishes 73(1):21–36 DOI 10.1007/s10641-004-3797-1.

Garvey JE, Herra TP, Leggett WC. 2002. Protracted reproduction in sunfish: the temporal dimension in fish recruitment revisited. Ecological Applications 12(1):194–205 DOI 10.1890/1051-0761(2002)012[0194:PRBFB]2.0.CO;2.

Garvey JE, Stein RA. 1998. Competition between larval fishes in reservoirs: The role of relative timing of appearance. Transactions of the American Fisheries Society 127(6):1021–1039 DOI 10.1577/1548-8659(1998)127<1021:CBLFIB>2.0.CO;2.
Gelwick FP, Matthews WJ. 1990. Temporal and spatial patterns in the littoral-zone fish assemblages of a reservoir (Lake Texoma, Oklahoma-Texas, USA). *Environmental Biology of Fishes* 27(2):107–120 DOI 10.1007/BF00001940.

Gido KB, Hargrave CW, Matthews WJ, Schnell GD, Pogue DW, Sewell GW. 2002. Structure of littoral-zone fish communities in relation to habitat, physical, and chemical gradients in a southern reservoir. *Environmental Biology of Fishes* 63(3):253–263 DOI 10.1023/A:1014359311188.

Hoxmeier JH, Aday DD, Wahl DH. 2009. Examining interpopulation variation in bluegill growth rates and size structure: effects of harvest, maturation, and environmental variables. *Transactions of the American Fisheries Society* 138(2):423–432 DOI 10.1577/T08-076.1.

Jackson JR, Noble RL. 2000. Relationships between annual variations in reservoir conditions and age-0 largemouth bass year-class strength. *Transactions of the American Fisheries Society* 129(3):699–715 DOI 10.1577/1548-8659(2000)129<0699:RBA VIR>2.3.CO;2.

Jackson ZJ, Quist MC, Larscheid JG. 2008. Growth standards for nine North American fish species. *Fisheries Management and Ecology* 15(2):107–118 DOI 10.1111/j.1365-2400.2007.00591.x.

Jolley JC, Edwards KR, Willis DW. 2009. Bluegill (*Lepomis macrochirus*) spawning periodicity and hatching duration in the Northern great plains, USA. *Journal of Freshwater Ecology* 24(1):29–38 DOI 10.1080/02705060.2009.9664262.

Kaemingk MA, Stahr KJ, Jolley JC, Holland RS, Willis DW. 2014. Evidence for bluegill spawning plasticity obtained by disentangling complex factors related to recruitment. *Canadian Journal of Fisheries and Aquatic Sciences* 71(1):93–105 DOI 10.1139/cjfas-2013-0282.

Kennedy RH, Walker WW. 1990. Reservoir nutrient dynamics. In: Thornton KW, Kimmel BL, Payne FE, eds. *Reservoir limnology: ecological perspectives*. New York, NY: John Wiley and Sons, 109–131.

Kimmel BL, Lind OT, Paulson LJ. 1990. Reservoir primary production. In: Thorton KW, Kimmel BL, Payne FE, eds. *Reservoir limnology: ecological perspectives*. New York, NY: John Wiley and Sons, 133–194.

Leonard DM, DeVries DR, Wright RA. 2010. Investigating interactions between channel catfish and other sport fishes in small impoundments. *North American Journal of Fisheries Management* 30(3):732–741 DOI 10.1577/M09-141.1.

Lienesch PW, Matthews MJ. 2000. Daily fish and zooplankton abundances in the littoral zone of Lake Texoma, Oklahoma-Texas, in relation to abiotic variables. *Environmental Biology of Fishes* 59(3):271–283 DOI 10.1023/A:1007688513523.

Michaletz PH. 2006. Prey resource use by bluegill and channel catfish in small impoundments. *Fisheries Management and Ecology* 13:347–354 DOI 10.1111/j.1365-2400.2006.00511.x.

Mittelbach GG. 1981. Foraging efficiency and body size—a study of optimal diet and habitat use by bluegills. *Ecology* 62(5):1370–1386 DOI 10.2307/1937300.

Mittelbach GG, Persson L. 1998. The ontogeny of piscivory and its ecological consequences. *Canadian Journal of Fisheries and Aquatic Sciences* 55(6):1454–1465 DOI 10.1139/f98-041.

Newbrey MG, Bozek MA, Jennings MJ, Cook JE. 2005. Branching complexity and morphological characteristics of coarse woody structure as lacustrine fish habitat. *Canadian Journal of Fisheries and Aquatic Sciences* 62(9):2110–2123 DOI 10.1139/f05-125.

Olson MH. 1996. Predator-prey interactions in size-structured fish communities: implications of prey growth. *Oecologia* 108(4):757–763 DOI 10.1007/BF00329052.
Partridge DG, DeVries DR. 1999. Regulation of growth and mortality in larval bluegills: implications for juvenile recruitment. *Transactions of the American Fisheries Society* 128(4):625–638 DOI 10.1577/1548-8659(1999)128<0625:ROGAMI>2.0.CO;2.

Ruhl N. 2013. The lotic-lentic gradient in reservoirs and estuaries. Ohio University Dissertation, pp. 12–21 and 76–94.

Santucci VJ, Wahl DH. 2003. The effects of growth, predation, and first-winter mortality on recruitment of bluegill cohorts. *Transactions of the American Fisheries Society* 132(2):346–360 DOI 10.1577/1548-8659(2003)132<0346:TEOGPA>2.0.CO;2.

Shoup DE, Callahan SP, Wahl DH, Peirce CL. 2007. Size-specific growth of bluegill, largemouth bass and channel catfish in relation to prey availability and limnological variables. *Journal of Fish Biology* 70:21–34 DOI 10.1111/j.1095-8649.2006.01204.x.

Stein RA, DeVries DR, Dettmers JM. 1995. Food-web regulation by a planktivore: exploring the generality of the trophic cascade hypothesis. *Canadian Journal of Fisheries and Aquatic Sciences* 52(11):2518–2526 DOI 10.1139/f95-842.

Theel HJ, Dibble ED. 2008. An experimental simulation of exotic aquatic macrophyte invasion and its influence on foraging behavior of bluegill. *Journal of Freshwater Ecology* 23(1):79–89 DOI 10.1080/02705060.2008.966459.

Thornton KW. 1990. Sedimentary processes. In: Thornton KW, Kimmel BL, Payne FE, eds. *Reservoir limnology: ecological perspectives*. New York, NY: John Wiley and Sons, 43–70.

Tomcko CM, Pierce RB. 2001. The relationship of bluegill growth, lake morphometry, and water quality in Minnesota. *Transactions of the American Fisheries Society* 130(2):317–321 DOI 10.1577/1548-8659(2001)130<0317:TROBGL>2.0.CO;2.

Tomcko CM, Pierce RB. 2005. Bluegill recruitment, growth, population size structure, and associated factors in Minnesota Lakes. *North American Journal of Fisheries Management* 25:171–179 DOI 10.1577/M04-054.1.

Vanni MJ, Arend KK, Bremigan MT, Bunnell DB, Garvey JE, Gonzalez MJ, Renwick WH, Soranno PA, Stein RA. 2005. Linking landscapes and food webs: effects of omnivorous fish and watersheds on reservoir ecosystems. *Bioscience* 55(2):155–167 DOI 10.1641/0006-3568(2005)055<0155:LLAFWE>2.0.CO;2.

Werner RG. 1969. Ecology of limnetic bluegill (*Lepomis macrochirus*) fry in Crane Lake Indiana. *American Midland Naturalist* 81(1):164–181 DOI 10.2307/2423658.

Werner EE, Hall DJ. 1979. Foraging efficiency and habitat switching in competing sunfishes. *Ecological Society of America* 60(2):256–264 DOI 10.2307/1937653.

Werner EE, Hall DJ. 1988. Ontogenetic habitat shifts in bluegill: the foraging rate-predation risk trade-off. *Ecological Society of America* 69(5):1352–1366 DOI 10.2307/1941633.

Werner EE, Hall DJ, Laughlin DR, Wagner DJ, Wilsmann LA, Funk FC. 1977. Habitat partitioning in a freshwater fish community. *Journal of the Fisheries Research Board of Canada* 43(3):360–370 DOI 10.1139/f77-058.

Werner EE, Peacor SD. 2003. A review of trait-mediated indirect interactions in ecological communities. *Ecology* 84(5):1083–1100 DOI 10.1890/0012-9658(2003)084[1083:AROTII]2.0.CO;2.

Wildhaber ML, Lamberson PJ. 2004. Importance of the habitat choice behavior assumed when modeling the effects of food and temperature on fish populations. *Ecological Modeling* 175:395–409 DOI 10.1016/j.ecolmodel.2003.08.022.