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FINAL REPORT SECTION 6 ENDANGERED SPECIES ACT



FEDERAL AID PROJECT E-12

HABITAT USE AND REPRODUCTIVE BIOLOGY OF ARKANSIA WHEELERI (MOLLUSCA: UNIONIDAE) IN THE KIAMICHI RIVER, OKLAHOMA

JUNE 1, 1990 - MAY 31, 1993

FINAL REPORT

STATE: Oklahoma

PROJECT NO: E-12

PROJECT TITLE:

Habitat use and reproductive biology of Arkansia wheeleri

(Mollusca: Unionidae) in the Kiamichi River, Oklahoma.

SEGMENT DATES: 1 JUNE 1990 - 31 MAY 1993

ABSTRACT

The only known remaining viable population of Arkansia wheeleri in the world occurs within an 80 mile stretch of the Kiamichi River in Pushmataha county, Oklahoma. Within this river A. wheeleri occurs only in the best available mussel habitat: backwaters and pools with fine gravel/coarse sand substrata, significant gravel bar and island development, emergent vegetation, and close proximity to riffles and tributaries. These areas harbor large, diverse mussel communities with which A. wheeleri is associated. In its optimal habitat, A. wheeleri is always rare: mean relative abundance varies from 0.2 to 0.7% and the average density is 0.27 individuals/m².

The reproductive biology and fish host(s) of A. wheeleri remain unknown. Two cyprinids, Notropis umbratilis and Notropis spp (c.f rubellus), are the most likely fish hosts. The youngest individual A. wheeleri encountered was approximately 12 years of age.

Forty-three percent of the historically known subpopulations of A. wheeleri below Jackfork Creek have apparently been extirpated and no new subpopulations have been located. A. wheeleri survive at 75% of the historically known locations above Jackfork Creek and five new subpopulations have been located. The relative abundance of A. wheeleri per site is slightly higher above Jackfork Creek than below. In addition, shell length data for live Amblema plicata, a dominant mussel species in the Kiamichi River, indicate reduced recruitment below Sardis Reservoir.

Much of the Kiamichi River watershed remains forested and this probably accounts for the high diversity and general health of its mussel community in comparison to other nearby rivers.

Program Narrative Objective

To determine the distribution, abundance and reproductive biology of the freshwater mussel Arkansia wheeleri within different habitats in the Kiamichi River of Oklahoma.

Job Procedures

- 1. Characterize microhabitats and determine the effects of impoundment.
- 2. Determine movement, growth, and survivorship of individuals.
- Identify glochidia and fish host.
- 4. Examine impact of Sardis Reservoir on the populations.
- Determine historic and current land use within the current range of Arkansia wheeleri in the Kiamichi River.

A. Introduction

Arkansia (syn. Arcidens) wheeleri, the Ouachita rock pocketbook, is a freshwater mussel. Originally named Arkansia wheeleri by Ortmann and Walker in 1912, Clarke (1981, 1985) recognized Arkansia as a subgenus of Arcidens. The

species is considered by Clarke to be distinct. However, Turgeon et al (1988) have continued to use the binomial Arkansia wheeleri.

The historical distribution of *Arkansia wheeleri* was in the Ouachita and Little rivers in Arkansas and the Kiamichi River in Oklahoma, all south-flowing rivers out of the Ouachita Mountains (Figure 1). A survey by Clarke (1987) indicated that the species is probably extirpated from the Ouachita River and severely depleted in the Little River. In 1992 and 1993, relict shells of *A. wheeleri* were found in the Little River in Oklahoma below Pine Creek Reservoir (Vaughn 1993a). No live *A. wheeleri* have been found in the Little River in Oklahoma as of August, 1993.

A. wheeleri was first reported from the Kiamichi River by Isely (1925). Since the construction of a dam and reservoir in the lower reaches of the Kiamichi in the 1970s, some of the backwater areas where it was known to occur have been destroyed (Valentine and Stansbery 1971), and connection with potential habitats on the Red River and other tributaries to it has been blocked. Clarke (1987) surveyed the Kiamichi River at 11 sites from just above Hugo reservoir upstream to near Albion. He located live specimens at six sites and empty shells at an additional two sites of the 11 sites sampled. A total of 10 live individuals was found along approximately 70 miles of river sampled, indicating a sparse but well distributed population in the river.

In 1988 and 1989 Mehlhop and Miller (1989) conducted a survey for A. wheeleri in the Kiamichi River from above Pine Creek to Whitesboro. Prior to this study, the habitat of A. wheeleri was reported to be backwater reaches of rivers where current is slow and where there are relatively non-shifting deposits of silt/mud and sand

(Wheeler 1918, Isely 1925, Clarke 1987, C.M. Mather, pers. comm.). Such areas tend to be in water that is shallow during late summer months when rainfall is minimal. In the Kiamichi River, these backwater areas usually are found adjacent to sand/gravel/cobble bars that either are scoured clean or support emergent aquatic vegetation, mainly *Justicia americana*. In preparation for their survey, Mehlhop and Miller (1989) examined aerial photographs of the river to determine probable backwater sites. In the 1989 survey, A. wheeleri were found at 13 sites. Population size was estimated to be just over 1000 individuals. Mehlhop and Miller found that the species was not restricted to backwater habitats with silty substrate as previously thought. They found that it occurs in pools with rock substrate as well, a more common habitat type in the Kiamichi River. This suggests that the species is more widely dispersed throughout the river than formerly thought and probably has greater reproductive potential. The Kiamichi River in Oklahoma supports the only remaining substantial population of this mussel in the world (Mehlhop and Miller 1989).

Based on the above information A. wheeleri was proposed for listing as a federal endangered species in July, 1990 (Federal Register 55(141):29865-29868).

Following a public comment period, the species was actually listed as endangered in October, 1991 (Federal Register 56(205):54950-54957).

The Kiamichi River is a major tributary of the Red River. It flows for a total of 169 miles through the southeastern Oklahoma counties of Leflore, Pushmataha, and Choctaw (OWRB 1990). The river flows across the Ridge and Valley Belt of the Ouachita Mountain geologic province and the Dissected Coastal Plain province (Curtis

and Ham 1979). The total watershed area is 1,830 square miles. Currently only the dam at Hugo, OK impounds the main river channel. However, a dam on Jackfork Creek in Pushmataha Co., a tributary of the Kiamichi, impounds Sardis Reservoir. The vegetation cover in the watershed can be described as a patchwork of forest made up of short-leaf and loblolly pine, mesic oak forests, and diverse bottomland habitats in various stages of maturity. Another large component of the watershed coverage is made up of pasture and other agricultural lands.

B. Methods

1. Characterize microhabitat and determine effects of impoundment

In 1990, 31 sites in the Kiamichi River between Antlers and Albion were examined for the presence of *Arkansia wheeleri*. Twenty two of these sites were judged to be appropriate for intensive surveying and habitat analysis. The results of the 1990 survey allowed us to determine that *A. wheeleri* occurs within select pools and backwaters in the Kiamichi River (Vaughn 1991). In 1991 we selected ten of these twenty two sites for intensive habitat analysis and population monitoring of *A. wheeleri* (Figure 2) (Vaughn and Pyron 1992). The ten sites were chosen to be as evenly distributed as possible along the Kiamichi River between Antlers and Albion but still be reasonably accessible and included all sites where *A. wheeleri* had been located by us in 1990 and some sites where it had been found historically (Mehlhop and Miller 1989, Clarke 1987).

Mussel surveys were conducted during July and August 1990, July and August 1991, and June - October 1992 using timed searches. Quadrat sampling techniques

were used in addition to timed searches in 1991 and 1992. Quadrat sampling was necessary in order to calculate mussel densities. However, it is difficult to find rare species such as A. wheeleri using quantitative techniques such as quadrat and transect sampling, therefore the more thorough timed searches also were conducted. Timed mussel surveys (timed to standardize sampling effort) were conducted by hand searching with the aid of SCUBA in deeper areas and by hand searches in shallow areas in the following manner: (1) an area was selected for surveying; (2) the entire area was searched by at least two people for one hour; (3) all mussels encountered were removed to shore; (4) all mussels were immediately identified and measured (total shell length); (5) mussels were put back in the water as close to where they were removed as possible. Quadrat sampling was done with quarter meter square PVC pipe quadrats. Quadrat sampling was done prior to timed searches. Fifteen random quadrats were sampled for each pool or backwater area. Quadrats were searched by hand, with the aid of SCUBA in deeper areas, until all mussels had been recovered to a depth of 15 cm. Individual mussels were measured and returned to the mussel bed as in timed searches.

At each site we characterized the substratum on the stream bottom and measured water depth, water temperature, current velocity, conductivity, dissolved oxygen, and pH. Current velocity was measured 10 cm above the stream bottom with a Marsh-McBirney model 201 portable flow meter. Conductivity and dissolved oxygen were measured with YSI meters. pH was measured with a Fisher Accumet portable pH meter. In 1990, water samples were taken for phosphate and nitrate.

Phosphate and nitrate analyses were performed by the Oklahoma State Department of Health. Three replicate substratum samples were collected at each site. These were brought back to the laboratory and allowed to dry. Samples were dry sieved, weighed, and individual proportions of samples assigned to the appropriate substrate size class (in mm) following. Hynes (1970, p. 24). Standard sieving techniques do not segregate particles greater than about 2 mm in diameter (i.e. gravel from pebble from cobble). To determine the proportion of fine gravel, coarse gravel, pebble, and cobble in samples we took the proportion of the sample greater than 2 mm in diameter and randomly measured the diameter of 100 particles in that subsample (Dunne and Leopold 1978).

We used a variety of statistical techniques to explore any relationship between

A. wheeleri distribution and abundance and measured habitat parameters.

Associations between A. wheeleri and other species of mussels as well as fish species were calculated using UGPMA Clustering (on relative abundance data), Spearman

Rank correlations (on relative abundance data) and the Jaccard Index (on presence/absence data) (Ludwig and Reynolds 1988). Habitat affinities were examined using three ordination techniques, Principal Components Analysis,

Reciprocal Averaging, and Canonical Correspondence Analysis.

Determine movement, growth, and survivorship of individuals.

All A. wheeleri found were measured using digital calipers (height, width, and length), and individually marked using numbered, laminated plastic fish tags. A. wheeleri were returned to the precise location from which they were captured.

To obtain additional information on A. wheeleri size and age distribution we measured shells of A. wheeleri that we found on the Kiamichi River between 1990 - 1992, that had been deposited in the Oklahoma Museum of Natural History, and that are in the collection of Dr. Charles M. Mather at the University of Science and Arts of Oklahoma. We then counted external annuli on the shells we had collected and those in the OMNH (McMahon 1991). We used the above data to calculate shell length, width and height vs. number of annuli regression lines. Shell height vs. number of annuli produced the best fit, and the resulting equation was used to predict the number of annuli for live mussels that had been measured in the field.

In early fall 1992 we brought four live A. wheeleri back to our laboratory for observation. The maximum number of A. wheeleri we were allowed to sacrifice or bring back to the laboratory under Vaughn's endangered species subpermit was five. These individuals were housed in a 626 liter Frigid Units artificial stream along with other mussel species that normally co-occur with A. wheeleri. The artificial stream was housed in an unheated room and stream temperatures approximated outdoor air temperatures. Mussels were fed Argent artificial plankton, a mixture of whole egg powder, egg yolk, shrimp meal, and fish meal which has been used to successfully raise freshwater prawn and larval finfish. A. wheeleri were checked daily for any material extruding from the siphon or shell. This material was examined under a dissecting microscope.

Identify glochidia and fish host.

We collected fish by seining (2 X 1.2 m high with 0.5 cm mesh, 6 X 1.2 m high

with 0.5 cm mesh) for 45 minutes per site, including all available habitats. We used experimental gill-nets on several dates to capture larger fish that are missed by seining. Fish collection dates were: 10 and 23 July 1990; 13 August 1991; 12 and 13 October 1991; 7 December 1991; 23 June 1992; 7, 8, 9, and 26 July 1992; 18 and 19 September 1992; and 12 October 1992. Fish were killed and preserved in 10% formalin and later transferred to 50% propanol before identification in the laboratory. Gills of individual fish were examined for glochidia under a dissecting microscope. We recorded the number of glochidia per individual fish. We removed all glochidia (except when glochidia were extremely numerous) and placed them in 50% propanol for future identification. Fish will be deposited in the Oklahoma Museum of Natural History.

Drift samples for planktonic glochidia were collected during the summer and fall of 1991 and 1992. Drift nets were run for two hours and four replicate samples were collected from each site. Samples were preserved in buffered formalin and later transferred to 70% ethanol. Glochidia were counted under a dissecting microscope using standard plankton counting techniques (Lind 1979). Glochidia samples are currently being stored in Dr. Vaughn's laboratory and are available for identification by appropriate experts.

We sacrificed some common, female mussels in the field and examined them for glochidia. These samples are also currently being stored in Dr. Vaughn's laboratory.

At the suggestion of Dr. Mark Gorden, Tennessee Technological University, and as permitted under Vaughn's endangered species subpermit, we placed live A.

wheeleri in the field inside plastic bags filled with river water. A. wheeleri were then held in these bags in the shade at stream side for several hours. This technique causes a build up of carbon dioxide which causes some mussels to release any glochidia they are harboring, but does not unduly stress the mussels (M.E. Gordon, pers. comm.).

4. Examine impact of Sardis Reservoir on the populations.

The impact of Sardis reservoir was examined by statistical comparison of the data obtained in procedures 1, 2 and 3 above and below the inflow from Sardis Reservoir via Jackfork Creek (see results section).

Determine historic and current land use within the current range of <u>Arkansia</u> wheeleri in the Kiamichi River.

We used spectral data collected from a satellite and completed an unsupervised classification of a portion of the data distinguishing several different landuse categories within the Kiamichi River watershed. Landuse information about the Kiamichi River watershed was extracted using a single, full-scene Landsat Thematic Mapper (TM) image. The image data were recorded on a series of tapes in 7 separate spectral bands. Each band represents the brightness value for each pixel in the image within specific wavelength intervals of the electromagnetic spectrum. The potential range of spectral values for each band can be found in Table 1. The acquisition date for this image was 13 July, 1992. This particular date was chosen because it provided a clear and almost cloud-free image occurring during the summer of 1992. These image data have a spatial resolution of 30 x 30 meters per pixel and cover a total area

approximately 180 km on a side. All image processing and classification was done using the ERDAS 7.5 software package on a personal computer.

Preclassification

We used bands 1, 4, and 5 without any additional enhancement in carrying out the classification. This band combination has been shown to provide the best differentiation of general landuse types for TM imagery (Chavez et al. 1982, 1984; Jensen 1986). Our final landuse classification was based on one developed by Anderson et al. (1976) for use with natural resource applications.

The portion of the image that includes the Kiamichi River watershed and the headwaters of the Little River watershed was selected and subsetted into a separate file for processing using a state watershed map as a guide for determining watershed boundaries. This image file was then divided into east and west portions of the watershed in order to reduce the size of the image files and to allow for easier manipulation within the computer environment. The final image and classification is a combination of these two files.

Classification

An unsupervised classification was completed for each of the two image files (see Jensen, 1986 for a detailed discussion of supervised and unsupervised classifications along with the different clustering algorithms). Unsupervised classifications are generally used when there is little a priori knowledge of the study

¹Current watershed maps were at too large a scale to allow us to separate out this small portion of the Little River watershed. However, including it did not affect our results.

area. We chose a clustering routine that uses a minimum spectral distance classification algorithm to assign each pixel a class value (1-27) that corresponds to an individual spectral class. We consistently used the default values calculated by ERDAS for the following clustering parameters: 1) N=27 - maximum number of clusters, 2) R=3.75 - minimum spectral distance between clusters, 3) C=5.25 - maximum cluster radius, 4) M=256 - number of points to process before merging clusters, and 4) T=1% - cluster elimination threshold. The initial classification identified 27 spectral classes. We manually compared each class to the original, unenhanced, 3-band image in order to group like-classes together and produce a classified image with three general categories: forest, cut forest, and water/cloud shadow. A fourth spectral class included all the clouds that were visible in the image. Clouds covered less than 1 % of the total image area.

The clustering routine did not do well at distinguishing urban areas due to their generally small size and the fact that the high resolution of the image allowed for multiple landuse types to be detected within an urban area (e.g. road, lawn, woodlot, etc). Because of this, we digitized the larger urban areas (population of 3,000 or more) and created a file that was later incorporated into the final classification. Roads also did not cluster independently. They consistently clustered with other surface features such as clearcuts, pasture, and clouds. This was due to their spectral similarity to these classes and the fact that pixels that included roads were often dominated by other landuse types (e.g. forest). For this reason, a second digitized file was created to include the primary roads that were visible on the image. This

digitized file was later incorporated into the final classification.

From the original classification we separated each of the first three general categories (forest, cut forest, water/cloud shadows) into their own data file so that each could be further clustered to provide greater detail in landuse recognition. Each of these classes was clustered with the same clustering algorithm used in the initial classification. This gave us at least six additional spectral classes for each general category that could then be compared to ancillary data in order to make decisions about which spectral classes should be combined and into which of the final landuse categories they belonged. Ancillary data consisted of black and white aerial photography (scale of 1" = 660") of Pushmataha and LeFlore counties provided by the Agricultural Stabilization and Conservation Service, and a general knowledge of the landscape. The acquisition date for photography from Pushmataha County was 1991 while Leflore County photography dated from 1978. The cloud class was already sufficiently identified so that no additional clustering was required.

An exception to the use of the 3-band combination for clustering occurred when we used only band 4 to carry out the second classification of the forest landuse class.

Band 4 alone can better distinguish conifers from deciduous trees (Jensen 1986) and was used to help distinguish three forest types: coniferous, deciduous, and mixed.

Historical Land Use

Significant land use changes in the Kiamichi watershed occurred in the first few decades of this century when the area was extensively logged. Unfortunately, we were unable to obtain aerial photographs from this era.

C. Results

1. Characterize microhabitats and determine the effects of impoundment.

The Kiamichi River contains an abundant and diverse mussel fauna with a high proportion of rare species. The fauna has changed little since originally described in the 1920's (Isely 1925). Mean total densities of mussels by site are shown in Figure 3. Mean densities of mussel species by site are shown in Table 2. Mean relative abundances of individual mussel species are shown in Figure 4.

Water quality data and substrate data are shown in Table 3. Phosphate, nitrate, and water quality data collected by the USGS are given in Appendix 1. Most measured water quality parameters did not vary significantly between sites. Current velocity, water depth, and substratum composition did vary between sites.

Arkansia wheeleri are extremely rare in the Kiamichi River. Figure 5 shows the number of A. wheeleri found at ten study sites on the Kiamichi River from 1989 - 1992 (sites 1 - 5) and 1990 - 1992 (sites 6 - 10). In this figure individual mussels found in any one year may represent recaptures from an earlier year and, therefore, numbers of mussels found should not be totalled over years for a site (see paragraph on recaptures below).

In 1990 two individual mussels were found, one at site 4 and one at site 10, which we were unsure of whether they were immature A. wheeleri or a pustule-less morph of Quadrula pustulosa. Juvenile A. wheeleri and some unusual morphs of Q. pustulosa can be very difficult to tell apart. The only way to be absolutely sure of the identification of these two individuals would have been to sacrifice them and examine

characteristics inside the shell, which we did not have permission to do in 1990. To be on the safe side, we assumed that these individuals were A. wheeleri and marked them. In 1992 we received permission to sacrifice up to five A. wheeleri. In 1992 we opened some mussels that were identical to the ones that we had been unsure about in 1990. The opened mussels were Q. pustulosa. We have therefore corrected the data to show that during our study no live A. wheeleri were found at either site 4 or 10.

In most cases A. wheeleri were located only through timed searches and did not occur in quadrat samples. Mean relative abundance of A. wheeleri at individual sites in 1990-92 is shown in Figure 6 and varied from 0.2% to 0.7%. In 1991 A. wheeleri occurred in quadrat samples at two sites, 6 and 7. This allowed us to calculate the density of A. wheeleri at these two sites. The density of A. wheeleri was 0.27 individuals per square meter at both of these sites.

Arkansia wheeleri only occurs in large mussel beds in association with other mussel species. Mussel sites or "beds" where A. wheeleri occur are more species-rich than other mussel beds that we sampled in the Kiamichi River (Figure 7, t=3.18, df=15, P=0.006 (data for 22 sites from 1990).

Using data from the twenty-two sites sampled in 1990, we used cluster analysis, Spearman Rank Correlation and the Jaccard Index to look for associations between A. wheeleri and other mussel species. The strongest association was between A. wheeleri and Quadrula quadrula. This positive association was found with Spearman Rank correlations (Table 4) and the Jaccard Index (Table 5). The next strongest

association was between A. wheeleri and Ellipsaria lineolata, found using both Spearman Rank correlations and the Jaccard Index.

We used several different ordination techniques to explore the habitat affinities of A. wheeleri. Data used in these ordinations included mussel relative abundances, fish abundances, quantitative habitat data, and coded habitat data. Sites were scored or coded as to whether they were a backwater or a pool (Htype), contained emergent vegetation, were within one fourth mile of a tributary entering the Kiamichi, contained gravel bar development, longitudinal position along the river and whether they were above or below Sardis Reservoir. Substrata data were coded as to the dominant substratum type for the 1990 data. Quantitative substratum data were used for 1991-1992.

Data for eight sites for which we had both fish and mussel abundance data were analyzed using Principal Components Analysis (PCA). Data used were means for 1991-1992. Arkansia wheeleri loaded positively on PCA axis 1 and was highly negatively correlated with PCA axis 2 (Figure 9, Table 6). In this analysis the amount of sand and fine gravel also loaded negatively on PCA axis 2. Therefore, from this analysis A. wheeleri would be predicted to be found in areas with more fine gravel and sand.

Reciprocal averaging produces simultaneous ordinations of sites and species allowing one to examine relationships between sites and species in one analysis (Ludwig and Reynolds 1988). RA was performed on the same data used for the PCA. Our ordination using RA separated species better than sites (Figure 10). Pearson

correlations with the resulting first and second RA axes produced only two biologically meaningful relationships (Table 7). Flow correlated with the first RA axis and Htype (habitat) correlated with second axis. Depth, conductivity, and substratum type were not strongly correlated with either RA axis. A. wheeleri came out along the middle of both axes and showed no distinct habitat preferences in this analysis.

Canonical correspondence analysis (CCA) is a direct gradient ordination technique that relates species abundances to measured variation in the environment (Ter Braak 1986; Taylor et al. 1992; Pyron and Taylor 1993). We used CCA to determine if any association exists between measured habitat variables and mussel relative abundances, including the relative abundance of A. wheeleri. CCA was performed on data for the 22 sites sampled in 1990 and on data from the ten sites sampled intensively in 1991.

The results of the CCA ordination for the 22 1990 sites are shown in Figure 11. In this analysis the first CCA axis accounted for 42.9% of the variance and the second CCA axis accounted for 23.6% of the variation. In Figure 11 the top graph shows the approximate centers of species distributions along the first two CCA axes, the middle graph shows the habitat vectors along those axes, and the bottom graph shows the positions of the 22 study sites along the CCA axes. Numbering of these sites does not correspond to numbering of the ten sites from 1991-1992. For example site three in this ordination is not the same as site three in Figure 12 (see below). In this ordination all of the habitat variables were clumped and intercorrelated and mussel species were also clumped. Sites, however, were well spread out. This ordination

mainly separated "good" from "poor" mussel habitat. The majority of mussel species were most abundant at those sites falling in the middle of the ordination. These sites had in common the following habitat characteristics: pools or backwaters, substratum composed predominately of coarse sand and fine gravel, proximity to a tributary, emergent vegetation, gravel bars, islands, and low flow.

The results of the CCA ordination for the ten 1991 sites are shown in Figure 12. A. wheeleri was not associated with any distinct habitat vector in this analysis, but neither were most mussel species sampled. It is important to point out that the ten study sites were chosen because they were pools or backwaters that contained large mussel beds that at least historically harbored A. wheeleri. By definition these habitats should be very similar to one another, and mussel species with the ability to survive in a variety of microhabitats within these pools and backwaters would not be expected to show a distinct habitat preference in this analysis. Mussel species that do occupy specific microhabitats within pools and backwaters did show a distinct habitat preference in the ordination. For example, Lampsilis teres prefers sandy substrate, Megalonaias nervosa only occurs in large downstream pools, and Villosa arkansasensis only occurs in cobble substrate (Oesch 1984). All of these associations were identified by the CCA ordination (Figure 12). A variable ("habitat") was used in the ordination to distinguish pools from backwaters. A. wheeleri showed no preference for pools versus backwaters.

2. Determine movement, growth, and survivorship of individuals.

In 1990 we marked and released at the point of capture nine A. wheeleri. In 1991 we recaptured and released at the point of capture nine A. wheeleri. In 1991 we recaptured only two marked individuals, although we found nine live individuals (Figure 5). Both recaptured A. wheeleri were found at site 3. Both of these individuals were found within one meter of where they were released in 1990. No other live A. wheeleri were found at site 3. In 1992 we recaptured the same two A. wheeleri at site 3 that we had recaptured in 1991. The individuals were within a few meters of where they had been released in 1991. The recaptured individuals had not grown discernably and changes (<1mm) are within the margin of error of our calipers (.1mm). No other marked A. wheeleri were recaptured in 1992.

Four individual A. wheeleri were brought back to the laboratory on September 20, 1992. A. wheeleri survived as well in the laboratory as any other mussel species we brought back, which included most non-rare species in the Kiamichi River, and survived better than some very abundant species such as Lampsilis ovata and Amblema plicata. A. wheeleri appeared to be doing very well for the first few months. They appeared well, were actively siphoning, and produced pseudofeces on a regular basis. The first A. wheeleri died in early December and the second followed in late December. At this time a large number of other species of mussels held in the laboratory also died. The third A. wheeleri died on January 11 and the last individual survived until March 8, 1993, or almost six months.

The size distribution (means for 1990 - 92) for A. wheeleri in the Kiamichi River is shown in Figure 13. Lengths of individual A. wheeleri captured in 1990 - 1992 at each site are shown in Figure 14. In Figure 14 each bar represents an individual mussel.

Lengths of spent shells in the OMNH and USAO collections were significantly different than lengths of live A. wheeleri in the Kiamichi River (Figure 15, t=1.9, df=78, P=0.03).

The resulting regression equation for number of annuli on shell height was Y = (-.483)X + 49.62 (n=24, R²=0.467, P < 0.05). Predicted ages based on number of annuli for live A. wheeleri from the Kiamichi River are shown in Figure 16. Predicted ages of spent shells vs. live A. wheeleri were not significantly different (t=-0.84, df=54, P=0.19).

The youngest predicted age for a live A. wheeleri was 12 years. Using this method none of the A. wheeleri we encountered on the Kiamichi River during our study were produced after Sardis reservoir was filled in 1983.

3. Identify glochidia and fish host.

We were not able to identify the glochidia or fish host of A. wheeleri. A. wheeleri examined in the field and held in the laboratory were not gravid and thus we were unable to obtain any glochidia either for identification or fish-host studies.

We were, however, able to obtain indirect information narrowing the field of potential fish hosts for A. wheeleri. Fish abundances and the numbers of glochidia found on fish are shown in Tables 8 and 9, respectively. Table 10 gives Spearman

Rank correlation coefficients of the association between fish and mussel abundance.

Significant correlations are shown in bold type. A. wheeleri was positively associated with several cyprinid species, notably Notropis spp. (formerly Notropis rubellus), N. umbratilis, N. volucellus, and a darter, Percina copelandi. No glochidia were found on P. copelandi or N. volucellus, but glochidia were found on N. rubellus and N. umbratilis from several sites (Table 9).

As discussed above, we performed a Principal Components Analysis (PCA) ordination of fish and mussel relative abundances. The relative abundance of A. wheeleri and N. umbratilis both had a strong negative loading on PCA axis 2 (Figure 9, Table 10).

Densities of drifting glochidia are shown in Figure 17 and discussed in more detail in the next section.

4. Examine impact of Sardis Reservoir on the populations.

A. wheeleri occurs both above and below the inflow to the Kiamichi River from Sardis Reservoir via Jackfork Creek. Of our ten study sites selected for detailed study, three were located above Sardis Reservoir and seven below (Figure 2). All of these sites historically harbored A. wheeleri. A. wheeleri was found during this study at all three sites (100%) above Sardis Reservoir. A. wheeleri was found at three of seven (43%) of the sites below the reservoir inflow. The relative abundance of A. wheeleri at sites above Sardis reservoir was on average greater than the relative abundance of A. wheeleri at sites below the reservoir (Figure 6), although these differences are not statistically significant.

The smallest live A. wheeleri was found at site 2 (above the reservoir) and the next smallest at site 7 (below the reservoir). However, if our predicted age distributions are correct, both of these individuals are older than Sardis reservoir which was filled in 1983.

Overall, mussel densities vary both above and below Sardis Reservoir (Figure 3). Relative abundances of most mussel species are not significantly different above and below the reservoir (Figures 18 and 19).

To determine the effects of Sardis Reservoir on the recruitment of mussels in the Kiamichi River we examined the size distribution of *Amblema plicata*. *A. plicata* is a generalist mussel species that is extremely abundant in the Kiamichi River and occurred at all of our sites. Many juvenile mussels are extremely difficult to identify to species, but juvenile *A. plicata* are readily identifiable and we knew we had seen and measured them in the Kiamichi River. We measured the shell lengths of 1435 live *A. plicata* in the Kiamichi River in 1991. Shell lengths of *A. plicata* from above Sardis reservoir were significantly different than shell lengths of *A. plicata* from below the reservoir (F=9.55, P=.01). Smaller *A. plicata* were much more common above Sardis Reservoir than below (Figure 20).

We examined mean densities of glochidia drifting in the water column and compared these to mean mussel densities for each sites. Mean densities of glochidia in the drift are lowest at sites 4 and 5, the sites directly below and closest to the inflow from Jackfork Creek. Mussel densities, however, are not low at these sites (Figure 21).

Determine historic and current land use within the current range of <u>Arkansia</u> wheeleri in the Kiamichi River.

Every attempt was made to compare 1992 satellite imagery and recent aerial photography with older aerial photography in order to determine land use changes in the Kiamichi watershed. We could not locate pre-logging or 1930's aerial photography for the Kiamichi watershed. It is possible that some older photographs may exist in the National Archives, but we did not have the time or financial resources to pursue this source. 1954 photography for the Kiamichi watershed was available from ASCS in Salt Lake City. We did not use these photographs for two reasons. (1) We did not think they were old enough to be very useful since most logging (the significant land use change) occurred long before 1954. (2) They did not appear to be useful enough to be worth the expense and the long wait. ASCS has a minimum 12 week turn around time, but on other projects we have found it to be much longer. Our impression from the satellite image, photography we have examined, talking to locals, and the considerable amount of time we have spent driving around Pushmataha County, is that there has been very little recent development in the watershed, other than the construction of the two reservoirs.

The final classification categories were developed using the Anderson et al.

(1976) classification scheme only as a guide. Final classification categories include: 1)

urban, 2) primary roads, 3) pasture/regrowth/cropland, 4) deciduous forest, 5)

coniferous forest, 6) mixed forest, 7) rivers/streams/woodlands, 8) reservoirs and, 9)

clouds/cloud shadows (Figs. 25,26) Cropland is a separate land class category in the

Anderson et al. classification scheme but because this cover type comprises such a small part of our study area (C. Vaughn and D. Certain, personal observation) and is spectrally similar to pasture we chose to combine the two. The regrowth category was the only category included in our classification but not used in the Anderson et al. classification. This category was included due to the occurrence of many forest clearcuts in early stages of regeneration. The obvious areas of regrowth after forest clearcutting were combined with pasture and cropland because of spectral similarities. Areas of regrowth were distinguished from mature forest based on spectral and geometric features that are obvious when the image is viewed. These regrowth areas have high brightness values and are of a regular geometric shape. They often stand out in landscape since they are often surrounded by mature forest (Figure 27). Some of Anderson et al.'s categories were not included since they do not occur within the Kiamichi watershed (e.g. tundra, perennial snow or ice).

The percent coverage of each landuse category in terms of the number of pixels included in that class is detailed in Table 12. The mixed forest category includes the largest number of pixels overall; however, the number of pixels included in each category is not a direct measure of the areal coverage of that category due to the two-dimensionality of the image data. Accurate aerial coverage estimates require including topographic relief parameters such as slope and elevation. The clouds/cloud shadows category is not a landuse class and was not included in these calculations.

Error Analysis

Classification accuracy for each landuse category (omitting clouds/cloud shadows category since it is not a landuse and because it covers less than 1 percent of the total area) was estimated by choosing at least 20 reference data sites for each category from aerial photographs and transposing the location of those sites onto USGS 7.5' topographic maps. The maps were then used to locate the reference sites on the unenhanced, 3-band image. Each site was represented by a single pixel which was described by its true landuse type as determined by aerial photography interpretation. Each reference pixel was compared to its' corresponding pixel in the classified image to estimate the accuracy of classification (Table 13). User's accuracy estimates the probability that a pixel classified in the image actually represents that category on the ground (Congalton 1991). The producer's accuracy gives the probability of a reference pixel being correctly classified. User's accuracy for this classification ranges from 28.3% (mixed forest) to 100.0% (rivers/streams/woodlands). Producer's accuracy ranges from 0.0% (primary roads) to 100.0% (urban). The overall accuracy for this classification is 53% if we include all final landuse categories. However, there was a large degree of error introduced into the classification when the primary roads class was overlayed onto the spectrally classified image. A close look at that class alone overlayed on the raw image shows a poor matchup of pixels representing the class in each image. This displacement error was caused by the transformation of the vector data in the digitized files to gridded data usable by the classified image. If the primary roads category is not used in the accuracy

assessment then overall accuracy increases to 60.0%.

An additional source of classification error is the effect of topographic relief on recorded brightness values. Topographic relief can either increase or decrease brightness values of pixels depending on factors such as the slope and aspect of the terrain, and the sun angle. Pixels occurring on slopes facing direct sunlight often have exaggerated brightness values. The opposite is true for pixels on the shaded side of slopes. It is not unusual for pixels in high sunlight or complete shade to have the maximum (255) or minimum (0) brightness value, essentially covering up any spectral information about landuse type that might otherwise be visible. Pixels on east-facing slopes in this data set appeared to have exaggerated brightness values while west-facing slopes and narrow-canyon walls were often hidden in shade.

D. Discussion

Microhabitat

Arkansia wheeleri occurs in both pools and backwaters in the Kiamichi River, not just backwaters as was previously believed. However, while pool and backwater habitats are common in the Kiamichi River, A. wheeleri only occurs in a select few of them. Pools and backwaters where A. wheeleri occur have in common an (1) abundant and diverse assemblage of mussels, (2) bottom substrata that are stable and contain adequate amounts of fine gravel/coarse sand, (3) low current (but not stagnant), (4) low siltation, and (5) proximity to tributaries, emergent vegetation, riffles and gravel bars.

Although pools and backwaters were considered different habitat types in this

study, in most cases they are tightly interconnected and share many characteristics in common. Backwater areas tend to be shallower and have finer substrata. As backwaters merge into the main river channel they turn into deeper pools with coarser substrata and slightly higher current velocity. As stated before, at our sites A. wheeleri occurred in both of these microhabitats. In addition we believe A. wheeleri moves back and forth between these habitats either voluntarily or through physical displacement of shifting sediments. As described in the Results section, individuals at site three that were repeatedly recaptured had not moved. However, at another site (site five) we found unmarked individuals in the backwater area only for two years (1990 and 1991), and then in the pool area alone in 1992. At this site the backwater and pool were interconnected. This site had undergone a great deal of sediment deposition during the high flow of spring 1992 and a great deal of the original backwater sediment was shifted to the pool area.

Recent studies addressing the substratum preferences of unionids have reached different conclusions and substratum preferences among unionids remain poorly understood. However, mussels are generally believed to be most successful in stable, sand-gravel mixtures and are generally absent from substrata with heavy silt loads (Cooper 1984, Salmon and Green 1983, Stern 1983, Way et al. 1990). Most unionid species can be found on a number of different substrata, but growth rates of individuals in each microhabitat can be quite different (Kat 1982, Hinch et al. 1989). Furthermore, many mussel species can occupy a wide range of habitats as a result of extensive larval dispersal over a heterogenous stream environment (Strayer 1981), but

growth and reproduction may be optimized only under the habitat conditions described above. As an example consider *Amblema plicata*, the clearly dominant mussel species in the Kiamichi River. This species occurred in every microhabitat we examined (pool, backwater, riffle, run) and at every site we examined. Its density, however, was not the same in all of these habitats. The greatest numbers of individuals were found in the large, diverse mussel beds where *A. wheeleri* also occurred. It is clearly able to "survive" in a large number of habitats, but its survival and growth is only optimized in "good" habitat (Strayer 1981).

The key to the distribution of A. wheeleri in the Kiamichi River is the presence of the large mussel beds where other mussel species thrive. These shoals represent optimal habitat for most mussel species, as evidenced by the large number of species and their high abundance. These shoals usually contain both pool and backwater areas, have significant gravel bar development with accompanying vegetation (dominated by Justicia americana), and are close to a tributary (usually within one quarter mile). Shoals are usually adjacent to a major riffle area, although they can be either up or downstream of the riffle.

While other mussel species may survive in less than optimum habitat, A.

wheeleri clearly cannot. They only survive in the best available habitat. Other studies have shown that these mainstream river shoals in shallower areas with slow, steady current and vegetation and coarse substrate are optimal habitat for lotic unionids because of minimal turbulence, low silt and steady food supply (Salmon and Green 1982).

In summary, A. wheeleri does not show a habitat preferences that is unique from other unionids in the Kiamichi River. However, A. wheeleri only occurs in the best available habitat for mussels.

Movement, Growth and Survival

Locomotory tendencies differ among different mussel species. For example,

Anodonta grandis migrate up and down with changes in water level (White 1979) and
in this way avoid stranding at low water. Other species such as Uniomerus

tetralasmus and the introduced Corbicula fluminea remain in position and suffer
prolonged exposure to air (McMahon 1991). Marked individual A. wheeleri in a
backwater area (site 3) did not move significantly from July 1990 to July 1992.

However, at another site (site 5) unmarked individuals moved from a backwater area
into the adjacent pool area. This movement was probably the result of physical
displacement of these individuals through sediment scour and redeposition.

In the majority of mussel species the greatest amount of growth occurs in the first few years of life. Shell growth rate then declines exponentially with age, although the rate of tissue biomass accumulation usually remains constant (McMahon 1991). Our examination of live A. wheeleri in the Kiamichi River and of relict A. wheeleri shells in the museum collections indicate that this growth pattern is also followed by A. wheeleri. Early annuli (those near the umbo) are much wider than later annuli near the edge of the shell.

Recruitment, growth and survival of mussels is often assessed by monitoring changes in density and size demography of natural populations (Payne and Miller

1989). We have no quantitative historical data on densities of A. wheeleri in the Kiamichi River or anywhere else. Past size distribution, however, can be assessed by examining the size distribution of relict shells. The size distribution of live A. wheeleri in the Kiamichi River is skewed to the left (Sokal and Rohlf 1981) (Figure 15) with more large individuals and fewer small individuals than one would expect with a statistically normal distribution. The size distribution of relict shells (Figure 15) follows a more normal distribution, with a greater proportion of smaller individuals than in the live population. Looking at these shell length data alone one would conclude that the size distribution of A. wheeleri in the Kiamichi River has changed over time and recruitment has decreased.

External annular rings have long been used to determine mussel age and growth rates. Recently this technique has been heavily criticized as being replete with problems (Downing et al. 1992). Natural erosion and corrosion of shells makes it difficult to distinguish true from false annuli. For example, false annuli can be formed by the incorporation of small substrate particles into mussel shells. It is difficult to count closely deposited growth lines near the margins of old shells. This produces an underestimate of shell age that becomes more erroneous with shell age. Downing et al. (1992) studied populations of *Lampsilis radiata* and *Anondonta grandis* in an oligotrophic lake. In these populations, many mussels showed no new external annuli at all, even several years after individual animals had been marked. They concluded that estimates of growth based on shell annuli consistently overestimated real shell growth. In addition, shell size and growth rates are linked to environmental conditions.

For example, some species form narrower shells in coarser substrates (Hinch et al. 1989) or grow faster in sand than in mud (Hinch et al. 1986).

As described earlier, we counted annuli on relict shells and used the resulting shell height-annuli regression equation to predict number of annuli for live A. wheeleri. As pointed out by the above discussion, this method should be assumed to have a large margin of error and probably also underestimates ages of A. wheeleri. Using this method, the youngest live A. wheeleri we encountered was approximately 12 years of age and there were not significant differences between predicted ages of live individuals versus relict shells. No juveniles were encountered.

Both types of data, shell-size distributions and ages predicted from external annuli, demonstrated that most A. wheeleri encountered in the Kiamichi River are old.

<u>Life History</u>

Because of its rarity, the reproductive biology of A. wheeleri remains unknown. Like other anodontines, it is probably bradytictic. The closest relative of A. wheeleri, Arcidens confragosus, becomes gravid in the fall and releases glochidia in the spring (Clarke 1981). We were unable to obtain any gravid A. wheeleri and thus obtained no glochidia. A. wheeleri glochidia are probably similar to other alasmidontine glochidia. Alasmidontine glochidia are asymmetrical and have a stylet covered with microstylets which facilitate attachment to the fish host. Glochidial releases are probably tied to natural water temperature changes in the spring and fall (Jirka and Neves 1992).

The fish host or hosts of A. wheeleri remain unknown. However, we have

identified strong possibilities for the fish host species. *A. wheeleri* was positively associated with several cyprinid species which were found to harbor glochidia.

*Notropis (=Lythrurus) umbratilis, the redfin shiner, inhabits "sluggish pools lined with water willows (Justicia americana) over gravel or sand substrates" (Robison and Buchanan 1988). This is the same habitat occupied by *A. wheeleri*. *N. umbratilis* is widespread in the Mississippi and Ohio valleys and in the southern Great Lakes tributaries as far north as western New York, southern Ontario, southern Michigan and Wisconsin, and southeastern Minnesota. It occurs south in the Mississippi valley to the Red River drainage but is uncommon in tributaries east of the Mississippi River. It occurs west to central Kansas and Oklahoma in the Missouri, Arkansas and Red River drainages.

Notropis spp. (c.f. rubellus) is a new species that is currently being described by Drs. Julian Humphries at Cornell University and Robert C. Cashner at the University of New Orleans. The species description will be published in the first issue of Copeia (No. 1) in 1994. The range of Notropis spp. is from the Blue River throughout the Little River drainage, and includes the Kiamichi River (R.C. Cashner, pers. comm.). The taxonomy of the species in the Ouachita River is unresolved (R.C. Cashner, pers. comm.).

Determining the reproductive biology of A. wheeleri will be extremely challenging. It takes an average of four to six hours to locate one individual A. wheeleri at a known location. They are extremely rare. Obtaining enough A. wheeleri to perform standard life history studies would probably destroy any remaining viability

of the existing population. For example, determining the age at which A. wheeleri achieves sexual maturity and the number of years gamete production continues would necessitate sacrificing many A. wheeleri.

Identification of the fish host might best be done by a molecular genetic approach (DNA fingerprinting). Such analyses are being used by other researchers to identify fish hosts of mussels (White 1993). The technique compares DNA obtained from glochidia found attached to fish to a battery of DNA's from adult mussels in the community. Even this approach would not guarantee identification of the fish host. Identification is contingent upon A. wheeleri still reproducing (unknown), fish being collected during the spawning season of A. wheeleri (unknown), and collection of the correct fish host (unknown). We have tissue of three A. wheeleri from the Kiamichi River stored in an ultracool freezer at the University of Oklahoma. This material could be made available to researchers for DNA fingerprinting once they work out the techniques on more common species. We also have preserved glochidia samples available for analysis.

Effects from Sardis Reservoir

It appears that historically A. wheeleri did equally well above and below Jackfork Creek (Clarke 1987). Historically, A. wheeleri occurred at at least seven sites between Clayton and Antlers. However, in five years of combined sampling effort by Mehlhop and Miller, 1988-1989, and ourselves, 1990-1992, we have only found three subpopulations of A. wheeleri below Jackfork Creek. Therefore, only three out of seven or 43% of the known subpopulations of A. wheeleri survive below Jackfork

Creek. In contrast, three out of four or 75% of the historical locations of *A. wheeleri* above Jackfork Creek have been confirmed and five new locations have been discovered (Mehlhop and Miller 1989, Vaughn 1991). The fourth historical location above Jackfork Creek has not been adequately surveyed and may well contain a subpopulation of *A. wheeleri*. No new locations have been discovered below Jackfork Creek despite intensive survey efforts. In addition, the relative abundance of *A. wheeleri* is slightly higher above Jackfork Creek than below. Unfortunately, we have no historical abundance data for *A. wheeleri* in the Kiamichi River.

Overall mussel densities vary both above and below Sardis Reservoir and relative abundances of most mussel species are not significantly different above and below the reservoir. However, in any mussel survey it is easier to find large adults than small, secretive juveniles. As shown above with the A. wheeleri data, most adult mussels were probably produced before the reservoir was filled. Therefore, a finding of no differences in relative abundances of adult mussels above and below the reservoir may actually be a reflection of habitat conditions before reservoir construction. To determine the effects of Sardis Reservoir on the recruitment of mussels in the Kiamichi River we examined the size distribution of Amblema plicata. A. plicata is a generalist mussel species that is extremely abundant in the Kiamichi River and occurred at all of our sites. Many juvenile mussels are extremely difficult to identify to species, but juvenile A. plicata are readily identifiable. Shell lengths of live

²Biologists that attempted to survey this site from a canoe were threatened by a person on shore with a firearm.

A. plicata from above Sardis reservoir were significantly different than shell lengths of live A. plicata from below the reservoir. These data indicate that recruitment of A. plicata is reduced below Sardis Reservoir. Smaller A. plicata were much more common above Sardis Reservoir than below. Because A. plicata is a common, tolerant species, any reductions in its recruitment may signify similar problems with most mussel species in the community.

Recently malacologists have voiced concerns that many North American unionid populations are composed of slowly dwindling numbers of long-lived adults destined for extirpation as pollution and other disturbances prevent juvenile recruitment to aging populations (McMahon 1991).

The lowest average number of glochidia found in the drift occurred at sites 4 and 5, the two sites below and closest to the confluence with Jackfork Creek.

To date we have found no water chemistry differences at sites above and below Sardis Reservoir. However, this study was designed to gather broad information on river habitats used by A. wheeleri and is not an intensive investigation of water quality dynamics in proximity to Jackfork Creek. Nevertheless, we have observed large physical differences in water level and flow regime fluctuations above and below Sardis Reservoir. For example, site 4 (Clayton) is almost directly below the confluence with Jackfork Creek. The measured summer flow rates at this site are typically much higher than the other sites because of water being released from the reservoir. Periodic scouring of substrata exposed to high flow velocities can remove both substrata and mussels and prevent their successful resettlement (Young and Williams,

1983; McMahon, 1991). When we visited site 7 during the summer of 1991 water levels had obviously just dropped drastically. Our evidence for this was the large number of small pools on gravel bars that harbored live but rapidly perishing fish and mussels. We counted over 100 stranded mussels at this site. Water level variation can have significant effects on mussel survival and may pose a significant threat to A. wheeleri at sites below the confluence of Jackfork Creek. Declining water levels expose relatively immotile mussels for weeks or months to air. It is doubtful that A. wheeleri can withstand such long air exposure, especially during the hot southeastern Oklahoma summer. Water temperature in some of the pools of stranded animals exceeded 35°C. Adult mussels are fairly sedentary in habit. While most species are adapted to seasonal changes in water levels and flow rates, they cannot move fast enough to respond to unpredictable and rapid changes in water level and flow rate. Historical and Current Land Use

The primary landuse type within the Kiamichi watershed appears to be that of a mixed forest type with an even larger portion of the watershed being covered by mature or near-mature forest of some type. However, this classification also shows how human development has been concentrated along and immediately adjacent to the river channel. This is not surprising given the rugged nature of the landscape outside of the river valley. However, difficulty of access has not completely deterred development in this area as can be seen by the occurrence of many forest cuts in various stages of regrowth. Overall, this watershed still maintains significant coverage by stands of mature forest, but much of this forest is likely to differ dramatically from

its' original state prior to being cut. The most significant recent land use change in the watershed is the construction of Hugo and Sardis reservoirs.

E. Threats

The greatest threats to the continued existence of A. wheeleri in the Kiamichi
River are land use changes. The most serious land use changes are further
impoundment of the river, water transfers, timber harvesting, and pollution from
agricultural and industrial development. A. wheeleri is also threatened by the invasion
of exotic bivalve species, particularly the zebra mussel, Dreissena polymopha.
Impoundment and Water Transfers

Rivers regulated by dams differ from free-flowing rivers in many ways and alteration in volume of flow and timing of discharge can seriously impact riverine fauna. Stream organisms, including mussels, have evolved in rivers that experience seasonal low-flow and high-flow periods (Meador and Matthews 1992). Fluctuating flows, especially if there will be lower flows for long periods to time, will result in the stranding of many mussels. Unlike fish species which can move rapidly in and out of microhabitats with changes in water levels, mussels move very slowly and are unable to respond to sudden drawdowns. Even if stranding doesn't actually kill a mussel, desiccation and thermal extremes will cause physiological stress and may reduce reproductive potential (McMahon 1991). We have already observed significant stranding of mussel individuals in the Kiamichi River below Sardis Reservoir (Vaughn and Pyron 1992).

Fluctuating flows also mean that transport of particulates will vary. Depending

on the flow schedule and the materials normally transported in the water column, there is the potential for loss of organics which are the food base for mussels.

Increased flows associated with river regulation have the potential to alter the distribution of sediment through scour, flushing, and deposition of newly eroded materials from the banks. Increased flows have the potential to activate the bed (i.e. actually cause the bottom of the river to move). Bedload movement will wreak havoc on the survival of many mussels, particularly juveniles (Young and Williams 1983). Erosion caused by increased flows at one location results in deposition of this material further downstream. This "zone of aggradation" results in an increased width/depth ration of that portion of the channel. As width/depth ratios increase the potential for bedload transport also increases. Thus, increased flows cause habitat loss through both sediment deposition and increased bed mobility.

Sediment deposition not only removes habitat, but also clogs mussel siphons (i.e. smothers them) and interferes with feeding and reproduction (Aldridge et al. 1987).

In the long term, higher base flow levels and shorter periods between peak flood periods will decrease habitat complexity by preventing the formation of islands, establishment of macrophyte beds etc... (Frissell 1986). Stabilized sediments, sand bars, and low flow areas, are all preferred unionid habitats (Hartfield and Ebert 1986, Payne and Miller 1989, Stern 1983, Way et al. 1990). It is around these "complex" areas that most mussel beds in the Kiamichi River, and indeed the highest diversity of stream fauna, are found.

Flow regulation not only has the potential to profoundly effect the stream fauna, but riparian fauna as well. Flood waters that normally recharge soils and aquifers may be rapidly exported downriver. Lowered water tables may cause shrinkage of the riparian corridor and shifts in terrestrial species composition (Allan and Flecker 1993, Smith et al. 1991).

Because of their dependence on the appropriate substrate and flow conditions, freshwater mussels, including *A. wheeleri*, are already naturally patchily distributed in rivers. Any further fragmentation, such as the construction of a reservoir, will act to increase patchiness and to increase the distance between patches. These effects may have major consequences for the metapopulation (i.e. local or subpopulations connected by infrequent dispersal) structure of *A. wheeleri* (Vaughn 1993b). As some subpopulations are eliminated and dispersal distances are increased between other subpopulations, demographic and genetic constraints will diminish the ability of this species to respond to even natural stochastic events much less human-induced environmental change (Wilcox 1986, Murphy et al. 1990).

Timber Harvesting

Timber harvesting operations can have significant effects on both stream water quantity and quality. The influence of catchment vegetation on stream discharge is dependent on a large number of variables, many of which are site-specific. However, in general, removal of forest vegetation increases stream runoff (Campbell and Doeg 1989) and leads to many of the effects of increased flows discussed above.

Road-building activities and low water crossings associated with logging can

lead to the development of "headcuts", or migrating knickpoints in the channel remote from areas of actual modification. Headcuts result in severe bank erosion, channel widening, and depth reduction and can have devastating effects on the mollusc fauna (Hartfield 1993).

Pollution

Their sedentary life style and filter-feeding habits make mussels especially vulnerable to chemical pollution events. Contaminants can destroy mussel populations directly by exerting toxic effects or indirectly by causing or contributing to the elimination of essential food organisms or host fish (Havlik and Marking 1987). To date, the Kiamichi River has remained relatively unpolluted, and this is one reason it maintains a generally healthy mussel fauna. Rivers near the Kiamichi, which have experienced more development, are rapidly losing their mussel faunas. For example, below the point where the Little River receives effluent from a paper mill, there have been massive mussel die offs (Vaughn 1993a).

Predation

Natural predation does not appear to be a threat to A. wheeleri in the Kiamichi River. Fresh shells found opened along the shore are predominately Corbicula (Vaughn and Pyron, pers. obs.). Corbicula have been shown to be the dominant prey of muskrats in other systems in which it has invaded (Neves and Odum 1989).

Exotic Species

Zebra mussels (*Dreissena polymorpha*) are now found in the Arkansas River system in Oklahoma. The high dispersal capabilities of this species make it highly probable that it will invade the Red River system in the near future (French 1990).

Invasion of the Kiamichi would most likely be from the two existing reservoirs, Sardis and Hugo, because this is where boats (with encrusted adults or water containing larvae) would be launched. The zebra mussel could then spread downstream from both reservoirs. Construction of the authorized Tuskahoma Reservoir would provide an additional entryway for zebra mussels into the Kiamichi. The exotic bivalve Corbicula fluminea may also pose a threat to A. wheeleri (Mehlhop and Miller 1989).

Commercial Harvest

At this time harvest of mussels from the Kiamichi River for commercial purposes is minimal. However, commercial harvest could pose a grave threat in the future as other more accessible rivers are depleted of their mussel fauna.

F. Summary and Conclusions

The only known remaining viable population of *Arkansia wheeleri* in the world occurs within an 80 mile stretch of the Kiamichi River in Pushmataha County,

Oklahoma. Within this river *A. wheeleri* occurs only in the best available mussel habitat: backwaters and pools with fine gravel/coarse sand substrata, significant gravel bar and island development, emergent vegetation, and close proximity to riffles and tributaries. These areas harbor large, diverse mussel communities with which *A. wheeleri* is associated. *A. wheeleri* never occur in riffles, runs, stagnant backwaters, silted-in areas or in pools and backwaters lacking the habitat characteristics described above. *A. wheeleri* never occurs as a single species assemblage or in small mussel

beds. In its optimal habitat, A. wheeleri is always rare: mean relative abundance varies from 0.2 to 0.7% and the average density is 0.27 individuals/m².

The reproductive biology and fish host(s) of A. wheeleri remain unknown. Two cyprinids, Notropis umbratilis and Notropis spp (c.f rubellus), are the most likely fish hosts. The youngest individual A. wheeleri encountered was approximately 12 years of age.

Forty three percent of the historically known subpopulations of A. wheeleri below Jackfork Creek have apparently been extirpated and no new subpopulations have been located. A. wheeleri survive at at least 75% of the historically known locations above Jackfork Creek and five new subpopulations have been located. The relative abundance of A. wheeleri per site is slightly higher above Jackfork Creek than below. In addition, shell length data for live Amblema plicata, a dominant mussel species in the Kiamichi River, indicate reduced recruitment below Sardis Reservoir.

Much of the Kiamichi River watershed remains forested and this probably accounts for the high diversity and general health of its mussel community in comparison to other nearby rivers.

G. RECOMMENDATIONS

1. The existing population of Arkansia wheeleri in the Kiamichi River should be monitored on a long-term basis. In addition, the entire mussel community in the Kiamichi River should be monitored for any changes in mussel abundance and size class structure. The ten sample locations that we established in this study would

provide good long-term monitoring sites. In addition to A. wheeleri, the Kiamichi
River contains other rare mussel species and our data indicate that recruitment below
Sardis Reservoir is decreasing. Monitoring these trends in a timely manner could well
prevent the listing of other mussel species in the Kiamichi system.

- Maintenance of the entire fish assemblage in the Kiamichi River is essential for the survival of A. wheeleri, especially until the fish host is positively identified. Fish populations should be monitored.
- 3. Continued efforts should be made to determine the reproductive biology and fish host(s) of Arkansia wheeleri. We have identified strong possibilities for the host fish species, but without further study the actual host(s) will remain unknown. DNA fingerprinting is a technique which holds much promise in this area. Thin-sectioning of existing A. wheeleri shells would provide more accurate information about the historical and current age structure of the Kiamichi population. Studies of the reproductive biology of surrogate species may prove useful and should be pursued, providing that criteria to select appropriate surrogates can be identified.
- Additional rivers should be surveyed for the presence of Arkansia wheeleri.
 Recently, A. wheeleri shells (but not live individuals) have been found in the Little River in Oklahoma and in two tributaries to the Red River in Texas.

5. Efforts should be made to deter further habitat alteration in the Kiamichi River watershed. In particular, no further reservoirs should be constructed and clearcutting should be discouraged. Private landowners should be encouraged to leave an intact riparian zone along the rivers edge. The authorized Tuskahoma Reservoir would inundate upper reaches of the river inhabited by A. wheeleri and affect the remaining population and its habitat downstream. It should not be built. The proposed addition of hydropower to Sardis Reservoir would detrimentally impact the A. wheeleri population below the reservoir. Alterations in the natural flow regime as a result of the proposed water transfer project have the potential to devastate A. wheeleri populations in the lower reaches of the river.

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Caryn C. Vaughn, Ph.D.

Mark Pyron David L. Certain

Oklahoma Natural Heritage Inventory Oklahoma Biological Survey University of Oklahoma

Date: /2/1/93 1993

Approved: Oklahoma Department of Wildlife Conservation

By: Harold Namminga

Federal Aid/Research Coordinator

Band	Spectral range (μm)
1	0.45-0.52 (blue)
2	0.52-0.60 (green)
3	0.63-0.69 (red)
4	0.76-0.90 (reflective-infrared)
5	1.55-1.75 (mid-infrared)
6	2.08-2.35 (mid-infrared)
7	10.4-12.5 (thermal infrared)

Table 1. Characteristics of spectral bands of Landsat
5 Thematic Mapper image.

Table 4. Spearman Rank correlations of mussel relative abundances. Correlations that are significant at P ≤ .05 are shown in bold type.

	A. ligamentina	A. plicata	A. wheeleri	E. lineolata	F. flava	L. radiata	L. ovata	L. teres	M. nervosa	O. reflexa
A. ligamentina	1									
A. plicata	-0.097	1								
A. wheeleri	0.157	0.219	1							
E. lineolata	0.59	0.096	0.37	1						
F. flava	-0.303	-0.249	-0.273	-0.271	1					
L. radiata	0.262	-0.147	0.109	0.516	-0.192	1				
L. ovata	-0.065	0.023	-0.007	0.153	-0.081	0.259	1			
L. teres	-0.219	0.184	-0.368	-0.08	0.357	0.015	-0.085	- 1		
M. nervosa	0.267	-0.024	0.423	0.31	-0.215	0.472	-0.25	-0.088	1	
O. reflexa	0.149	-0.005	0.351	0.145	0.18	0.026	-0.484	-0.15	0.269	
O. jacksoniana	0.354	-0.073	0.082	0.518	-0.13	0.532	0.229	-0.254	0.392	-0.196
P. purpuratus	0.43	0.084	-0.03	0.56	-0.042	0.3	0.039	-0.137	-0.015	0.19
P. occidentalis	0.389	-0.059	0.167	0.36	-0.028	0.197	-0.016	-0.361	0.115	0.34
Q. pustulosa	-0.07	-0.656	0.085	-0.248	0.117	-0.042	-0.089	-0.518	0.148	0.229
Q. quadrula	0.138	-0.154	0.437	0.052	-0.086	0.208	0.086	-0.497	0.338	0.212
T. verrucosa	0.077	0.244	-0.169	0.066	0.141	-0.206	0.14	0.468	-0.153	-0.124
T. truncata	0.019	0.006	0.151	0.338	0.348	0.099	-0.083	0.14	-0.172	0.228
V. arkansasensis	0.016	0.27	-0.081	0.08	0.264	-0.185	-0.241	0.527	-0.034	0.333
V. lienosa	-0.063	0.1	-0.167	-0.014	0.403	0.005	0.286	0.43	-0.11	-0.171

Table 4. Spearman Rank correlations of mussel relative abundances. Correlations that are signficant at P ≤ .05 are shown in bold type.

O. jacksoniana	P. purpuratus	P. occidentalis	Q. pustulosa	Q. quadrula	T. verrucosa	T. truncata	V. arkansasensis	V. lienosa
1								
0.262	1							
0.135	0.429	1						
-0.006	0.025	0.205	1					
0.354	-0.021	0.336	0.487	1				
-0.336	-0.318	0.096	-0.425	-0.267	1			
0.095	0.284	0.314	-0.325	-0.226	0.052	1		NE T
-0.145	-0.11	0.041	-0.311	-0.207	0.353	0.27	1	
-0.012	-0.074	0.215	-0.146	-0.088	0.489	0.235	0.437	

Table 3. Mean physical and chemical habitat parameters for the ten sites.

	1	2	3	4	5	6	7	8	9	10
Average Flow (cm/s)	6	6	3.5	20	10	4	4	4	8	3
pH	7.3	7	7	7	7.4	7.6	7.6	7.1	7	7
Water temperature (oC)	29	29	26	29	30	28	28	30	30	28
Average Depth (cm)	29	91	55	59	79	120	80	76	45	110
Conductivity (umho/cm2)	44	52	47	50	50	52	53	55	40	40
Dissolved Oxygen (mg/l)	6.3	7	6.7	6.2	6.7	. 7	6.8	6.6	7	7
% Sand	9	9.3	25.6	8.5	4.6	16.8	11.3	7.8	20.7	9
% Fine Gravel	26.4	36	37.7	39.6	34	34.8	34.8	43	25.5	35.5
% Coarse Gravel	33.6	32	19.8	33.8	31.3	30.9	30.9	37	32.3	29.1
%Pebble	28	18.2	15.7	17.5	26.6	16.2	18.2	11	19.9	20.8
%Cobble	3.1	4.4	1.2	0.6	3.4	1.3	1.3	1.2	1.6	5.6

Table 2. Densities of mussel species (No./M2) by site. Species which were found during timed samples but didn't occur in quadrats are not shown. Data are means for 1991 - 1992.

	1	2	3	4	5	6	7	8	9	10
Actinonaias ligamentina	6.67	0.80	6.93	12.40	2.83	4.93	9.87	5.30	0.80	3.20
Amblema plicata	19.47	7.47	3.07	8.00	13.13	5.47	26.40	12.30	1.07	5.33
Arkansia wheeleri	0.00	0.00	0.00	0.00	0.00	0.27	0.27	0.00	0.00	0.00
Ellipsaria lineolata	0.00	0.00	0.00	0.80	0.00	0.40	1.60	12.27	0.00	0.27
Fusconaia flava	2.53	0.40	0.00	1.33	1.30	1.47	2.13	0.00	0.27	1.60
Lampsilis ovata	1.73	1.20	1.33	1.73	1.70	2.40	0.80	0.00	0.00	1.33
Lampsilis radiata	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.27	0.00	0.00
Lampsilis teres	1.33	0.00	0.13	0.00	0.13	0.27	0.00	0.27	0.00	0.00
Megalonaias nervosa	0.13	0.00	0.00	0.00	0.00	0.00	0.67	0.80	0.00	0.00
Obliquaria reflexa	0.00	0.13	0.53	0.27	0.50	0.40	0.27	0.00	0.53	0.40
Obovaria jacksoniana	2.67	0.13	0.13	0.13	0.30	0.00	0.13	0.27	0.00	0.27
Potamilus purpuratus	0.00	0.40	0.53	0.27	0.00	0.27	0.00	0.00	0.00	0.00
Ptychobranchus occidentalis	0.13	0.13	1.87	0.53	0.00	0.00	0.00	1.33	0.00	0.00
Quadrula pustulosa	2.80	4.67	9.73	7.47	6.77	12.00	5.47	5.87	16.80	11.87
Quadrula quadrula	0.80	0.27	0.40	0.40	0.13	0.13	0.13	0.27	0.00	0.67
Tritogonia verrucosa	2.67	1.07	2.00	0.67	0.13	0.00	0.93	1.60	0.00	0.80
Truncilia truncata	0.40	0.00	0.53	0.27	0.13	0.40	0.67	0.00	0.27	0.27
Villosa lienosa	0.13	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.13

Table 5. Results of Jaccard Index for Mussel Presence/Absence

		JI		JI
A. ligamentina	A. plicata	0.773 E. lineolata	F. flava	0.4
A. ligamentina	A. wheeleri	0.353 E. lineolata	L. radiata	0.5
A. ligamentina	E. lineolata	0.471 E. lineolata	L. ovata	0.421
A. ligamentina	F, flava	0.762 E. lineolata	L teres	0.25
A. ligamentina	L. radiata	0.429 E. lineolata	M. nervosa	0.4
A. ligamentina	L. ovata	0.714 E. lineolata	O, reflexa	0.429
A. ligamentina	L. teres	0.263 E. lineolata	O. jacksoniana	0.545
A. ligamentina	M. nervosa	0.353 E. lineolata	P. purpuratus	0.571
A. ligamentina	O. reflexa	0.444 E. lineolata	P. occidentalis	0.462
A. ligamentina	O. jacksoniana	0.611 E. lineolata	Q. pustulosa	0.364
A. ligamentina	P. purpuratus	0.632 E. lineolata	Q. quadrula	0.462
A. ligamentina	P. occidentalis	0.556 E. lineolata	T. verrucosa	0.4
A. ligamentina	Q. pustulosa	0.773 E. lineolata	T. truncata	0.571
A. ligamentina	Q. quadrula	0.556 E. lineolata	V. arkansasensis	0.417
A. ligamentina	T. verrucosa	0.85 E. lineolata	V. lienosa	0.2
A. ligamentina	T. truncata	0.632 F. flava	L. radiata	0.65
A. ligamentina	V. arkansasensis	0.176 F. flava	L. ovata	0.773
A. ligamentina	V. lienosa	0.167 F. flava	L teres	0.35
A. plicata	A. wheeleri	0.273 F. flava	M. nervosa	0.3
A. plicata	E. lineolata	0.364 F. flava	O, reflexa	0.6
A. plicata	F. flava	0.909 F. flava	O. jacksoniana	0.45
A. plicata	L. radiata	0.591 F. flava	P. purpuratus	0.7
A. plicata	L ovata	0.864 F. flava	P. occidentalis	0.55
A. plicata	L teres	0.318 F. flava	Q. pustulosa	0,909
A. plicata	M. nervosa	0.273 F. flava	Q. quadrula	0.55
A. plicata	O. reflexa	0.545 F. flava	T. verrucosa	0.818
A. plicata	O. jacksoniana	0.409 F. flava	T. truncata	0.619
A. plicata	P. purpuratus	0.636 F. flava	V. arkansasensis	0.15
A. plicata	P. occidentalis	0.5 F. flava	V. lienosa	0.2
A. plicata	Q. pustulosa	1 L. radiata	L. ovata	0.6
A. plicata	Q. quadrula	0.5 L. radiata	L. teres	0.25
A. plicata	T. verrucosa	0.909 L. radiata	M. nervosa	0.357
A. plicata	T. truncata	0,636 L radiata	O. reflexa	0.471
A. plicata	V. arkansasensis	0.136 L. radiata	O. jacksoniana	0.467
A. plicata	V. lienosa	0.182 L. radiata	P. purpuratus	0.588
A. wheeleri	E. lineolata	0.556 L. radiata	P. occidentalis	0.500
A. wheeleri	F. flava	0.238 L. radiata	Q. pustulosa	0.591
A. wheeleri	L. radiata	0.357 L. radiata	Q. quadrula	0.55
A. wheeleri	L. ovata	0.316 L radiata	T. verrucosa	0.5
A. wheeleri	L. teres	0.083 L. radiata	T. truncata	0.5
A. wheeleri	M. nervosa	0.333 L. radiata	V. arkansasensis	0.067
4. wheeleri	O. reflexa	0.385 L. radiata	V. lienosa	0.133
A. wheeleri	O. jacksoniana	0.364 L ovata	The state of the s	
Control of the Contro	P. purpuratus	0.333 L ovata	L. teres	0.238
A. wheeleri A. wheeleri	P. occidentalis	Charles and August on the August State of the Control of the Contr	M, nervosa	0.316
CONTRACTOR STATE OF THE PARTY O	A STATE OF THE PARTY OF T	0.308 L ovata	O. reflexa	0.471
A. wheeleri	Q. pustulosa	0.273 L ovata	O. jacksoniana	0.474
A. wheeleri	Q. quadrula	0.417 L. ovata	P. purpuratus	0.65
A. wheeleri	T. verrucosa	0.3 L ovata	P. occidentalis	0.5
4. wheeleri 4. wheeleri	T. truncata V. arkansasensis	0.429 L. ovata 0.125 L. ovata	Q. pustulosa Q. quadrula	0.864

Table 5. Results of Jaccard Index for Mussel Presence/Absence

A. wheeleri	V. lienosa	0.111 L. ovata	T. verrucosa	0.857
L. teres	M. nervosa	0.182 L. ovata	T. truncata	0.571
L. teres	O. reflexa	0.188 L. ovata	V. arkansasensis	0.1
L teres	O. jacksoniana	0.143 L. ovata	V. lienosa	0.211
L. teres	P. purpuratus	0.235 M. nervosa	O. reflexa	0.385
L. teres	P. occidentalis	0.125 M. nervosa	O. jacksoniana	0.5
L. teres	Q. pustulosa	0.318 M. nervosa	P. purpuratus	0.333
L. teres	Q. quadrula	0.125 M. nervosa	P. occidentalis	0.417
L. teres	T. verrucosa	0.35 M. nervosa	Q. pustulosa	0.273
L. teres	T. truncata	0.313 M. nervosa	Q. quadrula	0.417
L. teres	V. arkansasensis	0.429 M. nervosa	T. verrucosa	0.3
L. teres	V. lienosa	0.375 M. nervosa	T. truncata	0.333
O. reflexa	O. jacksoniana	0.313 M. nervosa	V. arkansasensis	0.125
O. reflexa	P. purpuratus	0.625 M. nervosa	V. lienosa	0.111
O. reflexa	P. occidentalis	0.643 O. jacksoniana	P. purpuratus	0.533
O. reflexa	Q. pustulosa	0.545 O. jacksoniana	P. occidentalis	0.429
O. reflexa	Q. quadrula	0.643 O. jacksoniana	Q. pustulosa	0.409
O. reflexa	T. verrucosa	0.524 O. jacksoniana	Q. quadrula	0.538
O. reflexa	T. truncata	0.529 O. jacksoniana	T. verrucosa	0.381
O. reflexa	V. arkansasensis	0.25 O. jacksoniana	T. truncata	0.533
O. reflexa	V. lienosa	0.143 O. jacksoniana	V. arkansasensis	0.091
P. purpuratus	P. occidentalis	0.667 O. jacksoniana	V. lienosa	0.182
P. purpuratus	Q. pustulosa	0.636 P. occidentalis	Q. pustulosa	0.5
P. purpuratus	Q. quadrula	0.563 P. occidentalis	Q. quadrula	0.692
P. purpuratus	T. verrucosa	0.619 P. occidentalis	T. verrucosa	0.476
P. purpuratus	T. truncata	0.647 P. occidentalis	T. truncata	0.563
P. purpuratus	V. arkansasensis	0.133 P. occidentalis	V. arkansasensis	0.167
P. purpuratus	V. lienosa	0.2 P. occidentalis	V. lienosa	0.25
Q. pustulosa	Q. quadrula	0.5 Q. quadrula	T. verrucosa	0.476
Q. pustulosa	T. verrucosa	0.909 Q. quadrula	T. truncata	0.471
Q. pustulosa	T. truncata	0.636 Q. quadrula	V. arkansasensis	0.167
Q. pustulosa	V. arkansasensis	0.136 Q. quadrula	V. lienosa	0.25
Q. pustulosa	V. lienosa	0.182 T. truncata	V. arkansasensis	0.214
T. verrucosa	T. trunçata	0.619 T. truncata	V. lienosa	0.286
T. verrucosa	V. arkansasensis	0.15 V. arkansasensis	V. lienosa	0.4

Table 6a. Principal component loadings for mussels.

Species	Mussel PC1	Mussel PC2	Mussel PC3	
V. lienosa	0.947	0.040	0.089	
V. arkansasensis	0.874	0.048	0.250	
T. verrucosa	0.788	0.245	0.250	
Q. quadrula	-0.788	0.209	0.456	
D. jacksoniana	0.724	0.483	0.432	
). reflexa	-0.721	0.556	0.177	
. teres	0.710	0.443	0.505	
. flava	0.658	0.518	-0.176	
). pustulosa	-0.589	0.376	0.590	
. ligamentina	-0.553	0.481	-0.385	
f. nervosa	-0.504	0.742	0.114	
radiata	0.150	0.898	-0.272	
truncata	-0.050	0.746	0.349	
wheeleri	0.164	-0.684	-0.246	
ovata	0.134	0.523	-0.772	
plicata	0.141	0.415	-0.783	
. lineolata	-0.401	0.354	-0.264	
occidentalis	-0.168	0.010	0.245	
purpuratus	-0.373	-0.485	0.156	

Table 6b. Principal component loadings from fish.

Species	Fish PC1	Fish PC2	Fish PC3
L. sicculus	-0.906	0.303	0.104
P. copelandi	0.862	-0.176	0.146
N. atherinoides	-0.850	0.324	0.192
C. whipplei	-0.831	0.308	-0.333
M. punctulatus	0.805	0.489	0.291
N. umbratilis	0.774	-0.541	0.017
E. radiosum	0.716	0.559	0.123
P. notatus	-0.687	-0.063	-0.335
F. notatus	0.557	0.617	0.333
P. vigilax	-0.522	-0.046	0.575
G. affinis	0.265	0.901	-0.183
L. megalotis	0.450	0.847	0.071
N. sutkussi	0.356	-0.824	-0.117
L. macrochirus	0.107	-0.818	-0.035
N. volucellus	0.412	-0.756	-0.137
F. olivaceus	-0.446	-0.609	0.402
L. cyanellus	0.015	0.059	0.844
N. boops	0.240	0.126	-0.627
P. sciera	0.082	0.469	-0.566
C. anomalum	0.478	-0.203	-0.240

Table 6c. Pearson correlation coefficients between environmental variables and principal components scores from fish and mussel PCA's.

Variable	Fish PC1	Fish PC2	Mussel PC1	Mussel PC2
Flour	0.070	0.000	0.450	0.050
Flow Depth	0.073 -0.247	-0.036 -0.183	-0.159 -0.322	-0.058 -0.104
Habitat	-0.247	0.470	0.053	0.562
Position	-0.775	-0.117	-0.768	0.462
Reservoir	-0.479	-0.163	-0.657	0.233
Sand	-0.602	0.273	-0.378	-0.179
Fine gravel	-0.407	-0.404	-0.557	-0.295
Coarse gravel	0.381	0.120	0.270	0.418
Pebble	0.738	-0.006	0.665	-0.017
Vegetation	0.647	0.652	0.856	0.322
Bars	0.280	0.307	0.362	0.176

	1st Axis	2nd Axis
Site 1	1.000	
Site 2	0.097	1.000
Flow	0.482	-0.218
Depth	-0.258	-0.065
Conductivity	-0.192	0.202
H Type	0.020	0.309
S Type	0.287	0.144

Table 7. Pearson correlations of habitat factors with Reciprocal Averaging Axes.

Table 8. Numbers of fish collected.

	SITE 1	SITE 1	SITE 2	SITE 3	SITE 3	SITE 3	SITE 4	SITE 4	SITE 4	SITE 4	SITE 5	SITE 6	SITE 7	SITE 7	SITE 7	SITE 7 Sep-92	SITE 10	SITE 10	SITE 10
	Jul-91	Jul-92	Jul-92	Oct-91	Dec-91	Jun-92	Aug-91	Oct-91	Dec-91	Jul-92	Jul-92	Sep-92	Jul-91	Oct-91	Dec-91	Sep-92	Oct-91	Dec-91	Oct-9
.episosteus platostomus													1						
ictiobus bubalus			1-2	-1															-
Noturus gyrinus								1					6	1					
Noturus noctumus											1.5	- 1				- 1			
Moxostoma erythrurum				- 1				di si	-0										
Dorosoma petenense											2	1					- 10		
Labidesthes sicculus	1	1	2	-33	- 4	3		33	3	9	. 3	8		68		7	60		
Minytrema melanops		1																	
Cyprinella venusta														2		- 4	7		
Cyprinella whippiei	1	2	- 4	- 25	26	15		49	16	26	- 6	13	- 1	107	10	11	50		
Opsopoedus emiliae				-										9					
Notropis atherinoides	34			28	137			- 61	33	1	3	21		68	14	15	127		
Notropis boops	2		. 3		6		4	9		31	- 4			30		1	7		
Notropis sp. (rubellus)		2	13		100	45	53	- 1		76	40	2		-		6	3		
Notropis umbratilis		- 5	16							3	24	1					- "		
Notropis valucellus			1			- 1		7	7	- 1	3		1-1-1		2				
Cempostome anomalum	1	2	13	5	3	- 8	- 6	- 4	9	8	- 1		1	22	3	- 1	3		
Pirnephales notatus	1	100	1	2	2	3		9	- 1	1		1		4	3	- 1	4	- 1	
Pimephales vigitax		1 18		3		-		6	- 5	- 1			1	1	- 1		5	2	
Gambusia affinis		4		3		- 1	- 1	-11		2		4	1	3	16	- 1			
Fundulus notatus	1					-	- 6						. 1				- 31		
Fundulus olivaceus	100		2	5		. 5		- 4		6	2			9	2	19	3		1
Micropterus punctulatus	2	- 1	1			- 1	- 4	- 1		2	1		. 1			- 6			
Micropterus salmoides									77			1					2		
Lepomis cyanellus		- 1	de la constante de la constant			1	- 1	2			2		1	1		2			1
Lepomis gulosus		- 0				1	13						1 2 4	1000			- V		
Lepomis macrochirus			4		- 3	- 1	18			17		- 1	.9	14	- 4	1	15		
Lepomis megalotis	- 1	. 1				. 9		39		14		5		7		9			
Pomoxis annularis										1	5			2					
Morone chrysops			4 -											1					
Crystallaria asprella																	- 4		
Etheostoma gracile																			
Etheostoma nigrum						- 15			2					2				2	
Etheostoma radiosum	8		- 6	- 5	- 4	3	1	26	7	12	100	2		2	2	1 1	5	1	
Percina caprodes								3	- "				3	1			1	7	
Percine copelandi		3	5		2	- 4		-1	2		. 5			1		2	2		
Percina phoxocephala				- 1	110												2		
Percina sciera	1	1	2	1						1		4		1	10	- 3			

Table 9. Glochidial infection rates for fish from the Kiamichi River.

	SITE 1	SITE 1	SITE 2	SITE 3	SITE 3	SITE 3	SITE 4	SITE 4	SITE 4	SITE 4	SITE 5	SITE 6	SITE 7	SITE 7	SITE 7	SITE 7	SITE 1	SITE 10	SITE 10)
and the second s	'Jul-91	'Jul-92	Jul-92	Oct-91	Dec-91	Jun-92	Aug-91	Oct-91	Dec-91	Jul-92	Jul-92	Sep-92	Jul-91	Oct-91	Dec-91	Sep-92	Oct-91	Dec-91	Oct-92	
Lepisosteus pletostomus	W. ASS		-	2000	2000000	Service Services	7.000		Parent Parent	10/07/09			- 1	1	20000	D DON'T LO	10000		2000	
Iceobus bubalus				0			- 1													5
Noturus gyrinus																				
Noturus noctumus				2.19							0	0				0			0	
Moxostoma erythrurum			1	0						1	F 10 1			1			-			
Dorosoma petenense							7				- 0	0							1000	
Labidesthes sicculus	0	0	0	0	. 0	0		0	- 1	0	0	0		0		0	0		0	
Minytrema melanops		0		77.00				Check	5 - 1	-				-			and the same		100	
Cyprinella venusta		100	5 1	1	100		34	15000		4-00	-			0	200		0.714		0	
Cyprinette whipplei	0	0.5	0	0.28	20-Jul	0.333		0.75	- 1	0.1	0.167	0.769	0	0.7	0.5	0.455	0.5		0.833	
Opsopoedus emiliae										1000				0						
Notropis atherinoides	0			0.2	0.35			0.75	- 1	0	Towns 1	0.381		0.5	0.214	0.6	0.6		0.25	
Notropis boops	0		0	112000	0			0	100	0.05	0.25	0		0.233		0	0.429		2000	
Notropis sp. (rubellus)			0.385		-	0.3	1	- 1		0.55	0.6	- 1		-		0.667	0			
Notropis umbratilis		0	0					100		0.3		0								
Notropis volucellus		5 00	0			- 1		0.286	. 1	0	0.3		2785		0					
Campostoma anomalum	0	0	0	0	0	0	0	all the ball of the same of	- 1	0	0		0.2	22	0	0	0			
Pimephales notatus			1	0.5	- 1	0		0.889	- 1	- 1		- 1	-	0.25	1	1	0.75	1		
Pimephales vigitax				0.3				0.667	1	- 1				0	- 1		0.6	1	0.6	
Gambusia affinis		0		0		0	0	0		0		0	0	0.3	0	0			0	
Fundulus notatus	0	1 00					0	110	5		-	- 1	0	1000	49	-36	0			5
Fundulus olivaceus			0	.0		0		0	4	0	0			0	0		0		0	2
Micropterus punctulatus	0	0	0			0	0	0	1	0	. 0		0							
Micropterus salmoides	1 2	100										0					0.5			
Lepomis cyanellus		0					0	0			0		0	0		0	C HAN		0	
Lepomis gulosus						0			3-3-1						100					
Lepomis macrochirus			0			0	0	0	4	0	0	0	0	0	0	0	- 0		0	2
Lepomis megalotis	0	0	0			0	-	0	1	0		0		0.143		0			0	1
Pomoxis annularis		15								0	0	-		0						
Morone chrysops													74	0						
Crystallaria esprella																	0	N.		
Etheostoma gracile										1	440								0	
Etheostoma nigrum				0					1		-			0	-		- 52			
Etheostome radiosum	0		0	0	0	0		0	1	0		0		0	0		0	0	0	
Percina caprodes								0					- 1	.0			0			
Percina copelandi		0	0		0	0		0	1		0			0		. 0	. 0			
Percina phoxocephala		-		0							- 8			25 1000			0		13	
Percina sciera	0		0	0				0	C I	0		0		0		0	0	2	0.	90

Table 10. Spearman Rank correlations of mussel vs. fish relative abundances. Figures shown in bold type are significant at P ≤ .05.

	L. sicculus	C. whipplei	N. atherinoides	N. boops	N. rubellus	N. umbratilis	N. volucellus	C. anomalum	P. notatus
A. ligamentina	0.643	0.714	0.548	0.253	0.048	-0.837	-0.101	-0.108	0.946
A. plicata	-0.048	-0.095	-0.429	0.193	-0.119	0.381	-0.127	0.347	-0.109
A. wheeleri	-0.376	-0.073	-0.097	-0.356	0.291	0.491	0.187	0.195	-0.204
E. lineolata	0.738	0.881	0.714	-0.157	-0.594	-0.545	-0.71	-0.393	0.182
F. flava	-0.263	-0.395	-0.527	0.236	-0.323	0.402	-0.147	0.271	-0.531
L. radiata	0.69	0.667	0.476	0.157	-0.857	-0.52	-0.862	-0.06	0.109
L. ovata	-0.286	-0.048	-0.524	0.868	0.214	0.038	0.254	0.719	0.473
L. teres	0.503	0.356	0.54	-0.36	-0.896	-0.268	-0.817	-0.167	-0.3
M. nervosa	0.873	0.655	0.518	-0.055	-0.546	-0.48	-0.697	-0.466	0.208
O. reflexa	0.814	0.599	0.659	0.055	-0.252	-0.868	-0.338	-0.542	0.549
O. jacksoniana	0.374	0.12	0.06	0.372	-0.699	-0.244	-0.578	0.042	-0.037
P. purpuratus	0.096	0.374	0.506	-0.36	0.337	-0.308	0.148	-0.624	0.037
P. occidentalis	0.323	0.419	0.563	0.042	-0.263	-0.727	-0.179	0.12	0.512
Q. pustulosa	0.905	0.762	0.929	-0.398	-0.476	-0.748	-0.621	-0.695	0.255
Q. quadrula	0.707	0.491	0.683	-0.188	0	-0.727	-0.14	-0.663	0.512
T. verrucosa	-0.357	-0.19	-0.214	0.157	-0.19	0.127	0.025	0.85	0.073
T. truncata	0.446	0.169	0.482	-0.268	-0.602	-0.558	-0.392	0.127	0.202
V. arkansasensis	-0.302	-0.206	-0.399	0.188	-0.385	0.63	-0.344	0.346	-0.567
V. lienosa	-0.764	-0.764	-0.791	0.124	-0.027	0.843	0.189	0.604	-0.667

Table 10. Spearman Rank correlations of mussel vs. fish relative abundances. Figures shown in bold type are significant at P ≤ .05.

P. vigilax	G. affinis	F. notatus	F. olivaceus	Micropterus	L. cyanellus	L. macrochirus	L. megalotus	E. radiosum	P. copelandi	P. sciera
0.464	0.084	-0.164	0.275	-0.561	-0.33	0.275		-0.311	-0.611	-0.156
-0.518	-0.323	-0.409	0.395	-0.122	0.368	0.395	-0.452	-0.372	0.503	0.012
-0.5	-0.25	-0.722	0.091	0.112	-0.562	-0.232	-0.521	0.116	0.482	0.159
-0.082	0.527	-0.3	-0.108	-0.781	0.101	-0.263	0.214	-0.443	-0.419	0
-0.439	0.084	0.302	-0.163	0.27	0.791	0.163	0.06	-0.127	0.47	-0.09
0	0.731	0.3	-0.192	-0.415	0.495	-0.371	0.595	-0.108	-0.263	0.108
-0.327	0.12	0.055	-0.156	0.39	-0.254	0.443	0.19	0.18	0.204	0.156
0.155	0.58	0.422	-0.117	-0.176	0.51	-0.827	0.479	0.21	-0.056	0.123
0.219	0.165	-0.094	0.288	-0.839	0.392	0.027	0.109	-0.48	-0.508	0.082
0.713	0.223	0.288	0.127	-0.65	0.179	0.12	0.371	-0.337	-0.855	-0.169
0.262	0.588	0.856	-0.261	0.111	0.488	-0.345	0.807	0.424	-0.261	0.382
0.028	0.182	-0.387	-0.388	-0.42	-0.411	-0.012	-0.036	-0.388	-0.503	-0.267
0.494	0.494	0.439	-0.145	0.049	-0.134	-0.422	0.539	0.277	-0.289	-0.193
0.546	0.323	0.027	0.12	-0.805	0.127	-0.347	0.262	-0.299	-0.731	-0.06
0.796	-0.036	0.082	0.265	-0.638	-0.121	0.072	0.132	-0.253	-0.855	-108
-0.191	0.156	0.218	0.012	0.659	-0.051	-0.395	0.119	0.551	0.683	-0.012
0.594	0.109	0.511	0.406	-0.049	0.578	-0.4	0.193	0.042	0.061	-0.424
-0.567	0.436	0.22	-0.463	0.493	-0.044	-0.581	0.385	0.726	0.394	0.788
-0.563	-0.096	0.219	-0.178	0.811	0.262	-0.137	-0.082	0.466	0.851	0.192

Table 11. Results of Jaccard Index for Fish and Mussel Presence/Absence.

Fish species	Mussel species	JI	Fish species	Mussel species	JI
Noturus noctumus	A. ligamentina	0.375	Gambusia affinis	A. ligamentina	0.625
Noturus nocturnus	A. plicata	0.375	Gambusia affinis	A. plicata	0.625
Noturus noctumus	A. wheeleri	0.25	Gambusia affinis	A. wheeleri	0.5
Noturus nocturnus	E. lineolata	0.375	Gambusia affinis	E. lineolata	0.625
Noturus noctumus	F. flava	0.375	Gambusia affinis	F. flava	0.625
Noturus noctumus	L. radiata	0.375	Gambusia affinis	L. radiata	0.625
Noturus noctumus	L. ovata	0.375	Gambusia affinis	L. ovata	0.625
Noturus noctumus	L. teres	0.25	Gambusia affinis	L. teres	0.5
Noturus noctumus	M. nervosa	0.25	Gambusia affinis	M. nervosa	0.25
Noturus noctumus	O. reflexa	0.375	Gambusia affinis	O. reflexa	0.625
Noturus noctumus	O. jacksoniana	0.375	Gambusia affinis	O. jacksoniana	0.625
Noturus noctumus	P. purpuratus	0.375	Gambusia affinis	P. purpuratus	0.625
Noturus noctumus	P. occidentalis	0.25	Gambusia affinis	P. occidentalis	0.625
Noturus noctumus	Q. pustulosa	0.375	Gambusia affinis	Q. pustulosa	0.625
Noturus noctumus	Q. quadrula	0.375	Gambusia affinis	Q. quadrula	0.625
Noturus noctumus	T. verrucosa	0.375	Gambusia affinis	T. verrucosa	0.625
Noturus noctumus	T. truncata	0.375	Gambusia affinis	T. truncata	0.625
Noturus noctumus	V. arkansasensis	0.125	Gambusia affinis	V. arkansasensis	0.25
Noturus noctumus	V. lienosa	0.125	Gambusia affinis	V. lienosa	0.125
Dorosoma petenense	A. ligamentina	0.25	Fundulus notatus	A. ligamentina	0.25
Dorosoma petenense	A. plicata	0.25	Fundulus notatus	A. plicata	0.25
Dorosoma petenense	A. wheeleri	0.25	Fundulus notatus	A. wheeleri	0.125
Dorosoma petenense	E. lineolata	0.25	Fundulus notatus	E. lineolata	0.25
Dorosoma petenense	F. flava	0.25	Fundulus notatus	F. flava	0.25
Dorosoma petenense	L radiata	0.25	Fundulus notatus	L. radiata	0.25
Dorosoma petenense	L. ovata	0.25	Fundulus notatus	L. ovata	0.25
Dorosoma petenense	L teres	0.125	Fundulus notatus	L teres	0.125
Dorosoma petenense	M. nervosa	0.125	Fundulus notatus	M. nervosa	0
Dorosoma petenense	O. reflexa	0.215	Fundulus notatus	O. reflexa	0.25
Dorosoma petenense	O. Jacksoniana	0.25	Fundulus notatus	O. jacksoniana	0.25
Dorosoma petenense	P. purpuratus	0.25	Fundulus notatus	P. purpuratus	0.25
Dorosoma petenense	P. occidentalis	0.125	Fundulus notatus	P. occidentalis	0.25
Dorosoma petenense	Q. pustulosa	0.25	Fundulus notatus	Q. pustulosa	0.25
Dorosoma petenense	Q. quadrula	0.25	Fundulus notatus	Q. quadrula	0.25
Dorosoma petenense	T. verrucosa	0.25	Fundulus notatus	T. verrucosa	0.25
Dorosoma petenense	T. truncata	0.25	Fundulus notatus	T. truncata	0.25
Dorosoma petenense	V. arkansasensis	0.125	Fundulus notatus	V. arkansasensis	in the first of the later of the later
Dorosoma petenense	V. lienosa	0.125	Fundulus notatus	V. lienosa	0.125
Labidesthes sicculus	A. ligamentina	1	Pimephales vigilax	A. ligamentina	0.375
Labidestries sicculus	A. plicata	1	Pimephales vigilax	A. plicata	0.375
Labidesthes sicculus	A. wheeleri	0.75	Pimephales vigilax	A. wheeleri	0.125
Labidesthes sicculus	E. lineolata	1	Pimephales vigilax	E. lineolata	0.375
Labidesthes sicculus	F. flava	1	Pimephales vigilax	F. flava	0.375
Labidesthes sicculus	L. radiata	0.875	Pimephales vigilax	L radiata	0.375
Labidesthes sicculus	L ovata	1	Pimephales vigilax	L. ovata	0.375
Labidesthes sicculus	L teres	0.625	Pimephales vigilax	L. teres	0.25
Labidesthes sicculus	M. nervosa	0.375	Pimephales vigilax	M. nervosa	0.125
Labidesthes sicculus	O. reflexa	1	Pimephales vigilax	O, reflexa	0.125
Labidesthes sicculus	O. jacksoniana	1	Pimephales vigilax	The state of the s	0.375
Labidesthes sicculus	P. purpuratus	1	Pimephales vigilax	O. jacksoniana P. purpuratus	0.375

Table 11. Results of Jaccard Index for Fish and Mussel Presence/Absence.

Labidesthes sicculus	P. occidentalis	0.75	Pimephales vigilax	P. occidentalis	0.375
Labidesthes sicculus	Q. pustulosa	1	Pimephales vigilax	Q. pustulosa	0.375
Labidesthes sicculus	Q. quadrula	1	Pimephales vigilax	Q. quadrula	0.375
Labidesthes sicculus	T. verrucosa	1	Pimephales vigilax	T. verrucosa	0.375
Labidesthes sicculus	T. truncata	1	Pimephales vigilax	T. truncata	0.375
Labidesthes sicculus	V. arkansasensis	0.375	Pimephales vigilax	V, arkansasensis	0
Labidesthes sicculus	V. lienosa	0.375	Pimephales vigilax	V. lienosa	0
Minytrema melanops	A. ligamentina	0.125	Fundulus olivaceous	A. ligamentina	0.75
Minytrema melanops	A. plicata	0.125	Fundulus olivaceous	A. plicata	0.75
Minytrema melanops	A. wheeleri	0.125	Fundulus olivaceous	A. wheeleri	0.5
Minytrema melanops	E. lineolata	0.125	Fundulus olivaceous	E. lineolata	0.75
Minytrema melanops	F. flava	0.125	Fundulus olivaceous	F. flava	0.75
Minytrema melanops	L. radiata	0.125	Fundulus olivaceous	L. radiata	0.625
Minytrema melanops	L. ovata	0.125	Fundulus olivaceous	L ovata	0.75
Minytrema melanops	L teres	0.125	Fundulus olivaceous	L. teres	0.375
Minytrema melanops	M. nervosa	0	Fundulus olivaceous	M. nervosa	0.25
Minytrema melanops	O. reflexa	0,125	Fundulus olivaceous	O. reflexa	0.75
Minytrema melanops	O. jacksoniana	0.125	Fundulus olivaceous	O. jacksoniana	0.75
Minytrema melanops	P. purpuratus	0.125	Fundulus olivaceous	P. purpuratus	0.75
Minytrema melanops	P. occidentalis	0.125	Fundulus olivaceous	P. occidentalis	0.5
Minytrema melanops	Q. pustulosa	0.125	Fundulus olivaceous	Q. pustulosa	0.75
Minytrema melanops	Q. quadrula	0.125	Fundulus olivaceous	Q. quadrula	0.75
Minytrema melanops	T. verrucosa	0.125	Fundulus olivaceous	T. verrucosa	0.7
Minytrema melanops	T. truncata	0.125	Fundulus olivaceous	T. truncata	0.7
Minytrema melanops	V. arkansasensis	0.125	Fundulus olivaceous	V. arkansasensis	-
Minytrema melanops	V. lienosa	0.125	Fundulus olivaceous	V. lienosa	0.2
Cyprinella whipplei	A. ligamentina	1	Micropterus punctulatus	A. ligamentina	0.5
Cyprinella whipplei	A. plicata	1	Micropterus punctulatus	A. plicata	0.5
Cyprinella whipplei	A. wheeleri	0.75	Micropterus punctulatus	A. wheeleri	0.37
Cyprinella whipplei	E. lineolata	1	Micropterus punctulatus	E. lineolata	0.
Cyprinella whipplei	F. flava	1	Micropterus punctulatus	F. flava	0.
Cyprinella whipplei	L. radiata	0.875	Micropterus punctulatus	L. radiata	0.37
Cyprinella whipplei	L. ovata	1	Micropterus punctulatus	The state of the s	0.
Cyprinella whipplei	L. teres	0.625		L. teres	0.12
Cyprinella whipplei	M. nervosa	0.375		M. nervosa	
Cyprinella whipplei	O. reflexa	1	THE RESIDENCE OF THE PARTY OF T	O. reflexa	0.
Cyprinella whipplei	O. jacksoniana	1	Micropterus punctulatus	The second secon	0.
Cyprinella whipplei	P. purpuratus	1	Micropterus punctulatus	AND THE PARTY OF T	0.
Cyprinella whipplei	P. occidentalis	0.75	NAME OF TAXABLE PARTY OF TAXABLE PARTY.	P. occidentalis	0.2
Cyprinella whipplei	Q. pustulosa	1			0.
Cyprinella whipplei	Q. quadrula	1	manifold and an expedimental and a contract through the form	THE RESERVE OF THE PARTY OF THE	0.
Cyprinella whipplei	T. verrucosa	1	Micropterus punctulatus	THE RESERVE OF THE PARTY OF THE	0.
Cyprinella whipplei	T. truncata	1	Micropterus punctulatus	and the body of the property of the contract o	0.12
Cyprinella whipplei	V. arkansasensis	0.375	Micropterus punctulatus		-
Cyprinella whipplei	V. lienosa	0.375	Micropterus punctulatus		0.12
Notropis atherinoides	A. ligamentina	0.875	Micropterus salmoides	A. ligamentina	0.2
Notropis atherinoides	A. plicata	1	Micropterus salmoides	A. plicata	0.2
Notropis atherinoides	A. wheeleri	0.625	Micropterus salmoides	A. wheeleri	0.12
Notropis atherinoides	E. lineolata	0.875	Micropterus salmoides	E. lineolata	0.2
Notropis atherinoides	F. flava	0.875	CONTRACTOR OF THE PROPERTY OF	F. flava	0.2
Notropis atherinoides	L. radiata	0.875		L. radiata	0.2

Table 11. Results of Jaccard Index for Fish and Mussel Presence/Absence.

L. ovata	0.875	Micropterus salmoides	L. ovata	0.25
L. teres	0.625	Micropterus salmoides	L. teres	0.25
M. nervosa	0.375	Micropterus salmoides	M. nervosa	0.25
O. reflexa	0.875	Micropterus salmoides	O. reflexa	0.25
O, jacksoniana	0.875	Micropterus salmoides	O. jacksoniana	0.25
P. purpuratus	0.875	Micropterus salmoides	P. purpuratus	0,25
P. occidentalis	0.75	Micropterus salmoides	P. occidentalis	0.25
Q. pustulosa	0.875	Micropterus salmoides	Q. pustulosa	0.25
Q. quadrula	0.875	Micropterus salmoides	Q. quadrula	0.25
T. verrucosa	0.875	Micropterus salmoides	T. verrucosa	0.25
T. truncata	0.875	Micropterus salmoides	T. trunçata	0.25
V. arkansasensis	0.25	Micropterus salmoides	V. arkansasensis	0.125
V. lienosa	0.25	Micropterus salmoides	V. lienosa	0
A. ligamentina	1	The state of the s	A. ligamentina	0.5
Control of the Contro	1			0.5
The second secon	0.75	THE RESERVE THE PROPERTY OF THE PARTY OF THE	A. wheeleri	0.375
E. lineolata	1	The state of the s	E. lineolata	0.5
	1	The state of the s		0.5
- Control of the Cont		and the second s	and the first backer be seen as a second	0.5
The second secon	1	The state of the s	The state of the s	0.5
THE RESERVE OF THE PARTY OF THE	-	CONTRACTOR OF THE PROPERTY OF	L. teres	0.375
The state of the s	-		M. nervosa	0.25
	_	The state of the s	- Service Control Control Control	0.5
A STATE OF THE PARTY OF THE PAR		The same of the sa	The second and the se	0.5
		THE RESERVE THE PROPERTY AND ADDRESS OF THE PARTY AND ADDRESS OF THE PA		0.5
- A STATE OF THE PARTY OF THE P	-	the state of the s	- participant was accommon	0.375
The second secon		Control of the Control of the Property of the Control of the Contr	CONTRACTOR OF STREET,	0.5
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August March Control of the Control		The second secon	THE RESERVE AND ADDRESS OF THE PARTY OF THE	0.75
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and the second distance of the second distanc	and the sales between the first	the state of the s	CONTRACTOR OF THE PARTY OF THE	0.625
The state of the s	THE RESIDENCE PROPERTY.		and the state of t	0.75
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	-		AND DESCRIPTION OF THE PROPERTY OF THE PROPERT	0.625
and the state of t	The second section and the	THE RESERVE AND ADDRESS OF THE PARTY OF THE	and the second and th	0.75
The second secon	- Anna	CONTRACTOR OF THE PARTY OF THE		0.375
The state of the s			and the same of the	0.375
THE RESIDENCE OF THE PARTY OF T		THE RESERVE OF THE PROPERTY OF		0.75
the state of the s	- International Control	The second contract of	Facilities of the Control of the Con	0.75
The second section of the second section of the second section			and the same of th	0.75
The state of the s	make the rate from the contract of the		THE PROPERTY OF THE PROPERTY O	0.75
and the second s	and a comment and committee had a	THE RESIDENCE OF THE PROPERTY	THE RESERVE AND PROPERTY OF THE PARTY OF THE	The second residence
AND DESCRIPTION OF THE PERSON NAMED IN COLUMN TWO IS NOT THE PERSON NAMED IN COLUMN TWO IS NAMED IN COLUM	and introduced by	The same of the sa	THE RESERVE AND ADDRESS OF THE PARTY OF THE	0.75
The state of the last of the state of the st	man and the second second second second	THE PROPERTY OF THE PROPERTY O	and the contract of the contra	0.75
	_		The second section is a second	0.75
		A SECURITION OF THE PROPERTY OF THE PARTY OF	The state of the s	40 56
T. truncata V. arkansasensis	0.875	Lepomis macrochirus Lepomis macrochirus	T. truncata V. arkansasensis	0.75
	L. teres M. nervosa O. reflexa O. jacksoniana P. purpuratus P. occidentalis Q. pustulosa Q. quadrula T. verrucosa T. truncata V. arkansasensis V. tienosa A. ligamentina A. plicata A. wheeleri	L. teres 0.625 M. nervosa 0.375 O. reflexa 0.875 O. jacksoniana 0.875 P. purpuratus 0.875 P. occidentalis 0.75 Q. pustulosa 0.875 Q. quadrula 0.875 T. verrucosa 0.875 T. truncata 0.875 V. arkansasensis 0.25 V. lienosa 0.25 A. ligamentina 1 A. wheeleri 0.75 E. lineolata 1 F. flava 1 L. radiata 0.875 L. ovata 1 L. teres 0.625 M. nervosa 0.375 O. reflexa 1 P. purpuratus 1 P. purpuratus 1 V. lienosa 1 J. truncata 1 V. lienosa 0.375 V. lienosa 0.375 V. lienosa 0.375 V. lienosa 0.375 A.	L. teres 0.625 Micropterus salmoides M. nervosa 0.375 Micropterus salmoides O. reflexa 0.875 Micropterus salmoides P. purpuratus 0.875 Micropterus salmoides P. purpuratus 0.875 Micropterus salmoides P. occidentalis 0.75 Micropterus salmoides Q. pustulosa 0.875 Micropterus salmoides Q. pustulosa 0.875 Micropterus salmoides Q. quadrula 0.875 Micropterus salmoides T. verrucosa 0.875 Micropterus salmoides T. verrucosa 0.875 Micropterus salmoides V. arkansasensis 0.25 Micropterus salmoides V. lienosa 0.25 Micropterus salmoides A. ligamentina 1 Lepomis cyanellus A. plicata 1 Lepomis cyanellus E. lineolata 1 Lepomis cyanellus E. lineolata 1 Lepomis cyanellus L. radiata 0.875 Lepomis cyanellus L. radiata 0.875 Lepomis cyanellus L. teres 0.825 Lepomis cyanellus D. purpuratus 1 Lepomis cyanellus P. occidentalis 0.75 Lepomis cyanellus P. occidentalis 0.75 Lepomis cyanellus V. arkansasensis 0.375 Lepomis macrochirus A. ligamentina 0.875 Lepomis macrochirus V. arkansasensis 0.375 Lepomis macrochirus V. arkansasensis 0.375 Lepomis macrochirus L. teres 0.825 Lepomis macrochirus L. Lepomis macrochirus D. picata 0.875 Lepomis macrochirus L. Lepomis macrochirus L. permis macrochirus D. purpuratus 0.875 Lepomis macrochirus L. permis macrochirus D. purpuratus 0.875 Lepomis macrochirus D. purpuratus 0.875 L	L. teres 0.625 Micropterus salmoides M. nervosa 0.375 Micropterus salmoides M. nervosa 0.75 Micropterus salmoides O. reflexa 0.875 Micropterus salmoides O. reflexa 0.875 Micropterus salmoides O. jacksoniana P. purpuratus 0.875 Micropterus salmoides P. purpuratus P. occidentalis 0.75 Micropterus salmoides P. purpuratus Q. pustulosa 0.875 Micropterus salmoides Q. pustulosa Q. quadrula 0.875 Micropterus salmoides Q. pustulosa Q. quadrula 0.875 Micropterus salmoides Q. quadrula T. verrucosa 0.875 Micropterus salmoides Q. quadrula 0.875 Micropterus salmoides Q. quadrula V. arkansasensis V. arkansasensis V. arkansasensis V. arkansasensis V. arkansasensis V. arkansasensis V. lienosa 0.25 Micropterus salmoides V. arkansasensis V. lienosa 1 Lepornis cyanellus A. figamentina A. plicata 1 Lepornis cyanellus A. plicata 1 Lepornis cyanellus E. lineolata 1 Lepornis cyanellus E. lineolata 1 Lepornis cyanellus E. lineolata 1 Lepornis cyanellus L. radiata 1 Lepornis cyanellus L. ovata 1 Lepornis cyanellus L. teres M. nervosa 0.375 Lepornis cyanellus D. reflexa 0. jacksoniana 1 Lepornis cyanellus P. purpuratus 1 Lepornis cyanellus P. purpuratus 1 Lepornis cyanellus P. purpuratus P. occidentalis 0.75 Lepornis cyanellus P. purpuratus P. occidentalis 0.75 Lepornis cyanellus P. poccidentalis Q. pustulosa 1 Lepornis cyanellus V. arkansasensis V. lienosa 0.375 Lepornis cyanellus T. truncata V. arkansasensis 0.375 Lepornis macrochirus A. ligamentina A. plicata 0.875 Lepornis macrochirus A. plicata 1. Lepornis macrochirus A. plicata 0.875 Lepornis macrochirus D. reflexa 0.875 Lepornis macrochirus D. purpuratus 0.975 Lepornis macrochirus D. purpuratus 0.975 Lepornis mac

Table 11. Results of Jaccard Index for Fish and Mussel Presence/Absence.

Notropis umbratilis	A. ligamentina	0.5	Lepomis megalotis	A. ligamentina	0.875
Notropis umbratilis	A. plicata	0.5	Lepomis megalotis	A. plicata	0.875
Notropis umbratilis	A. wheeleri	0.375	Lepomis megalotis	A. wheeleri	0.75
Notropis umbratilis	E. lineolata	0.5	Lepomis megalotis	E. lineolata	0.875
Notropis umbratilis	F. flava	0.5	Lepomis megalotis	F. flava	0.875
Notropis umbratilis	L. radiata	0.375	Lepomis megalotis	L. radiata	0.75
Notropis umbratilis	L. ovata	0.5	Lepomis megalotis	L ovata	0.875
Notropis umbratilis	L. teres	0.25	Lepomis megalotis	L. teres	0.625
Notropis umbratilis	M. nervosa	0.125	Lepomis megalotis	M. nervosa	0.375
Notropis umbratilis	O. reflexa	0.5	Lepomis megalotis	O. reflexa	0.875
Notropis umbratilis	O. jacksoniana	0.5	Lepomis megalotis	O. jacksoniana	0.875
Notropis umbratilis	P. purpuratus	0.5	Lepomis megalotis	P. purpuratus	0.875
Notropis umbratilis	P. occidentalis	0.25	Lepomis megalotis	P. occidentalis	0.75
Notropis umbratilis	Q. pustulosa	0.5	Lepomis megalotis	Q. pustulosa	0.875
Notropis umbratilis	Q. quadrula	0.5	Lepomis megalotis	Q. quadrula	0.875
Notropis umbratilis	T. verrucosa	0.5	Lepomis megalotis	T. verrucosa	0.875
Notropis umbratilis	T. truncata	0.5	Lepomis megalotis	T. truncata	0.875
Notropis umbratilis	V. arkansasensis	0.375	Lepomis megalotis	V. arkansasensis	estation productions
Notropis umbratilis	V. lienosa	0.375	Lepomis megalotis	V. lienosa	0.25
Notropis valucellus	A. ligamentina	0.375	Etheostoma radiosum	A. ligamentina	0.875
Notropis volucellus	A. plicata	0.375	Etheostoma radiosum	A. plicata	0.875
Notropis volucellus	A. wheeleri	0.25	Etheostoma radiosum	A. wheeleri	0.625
Notropis volucellus	E. lineolata	0.375	Etheostoma radiosum	E. lineolata	0.875
Notropis volucellus	F. flava	0.375	Etheostoma radiosum	F. flava	0.875
Notropis volucellus	L. radiata	0.25	Etheostoma radiosum	L. radiata	0.75
Notropis volucellus	L. ovata	0.375	Etheostoma radiosum	L. ovata	0.875
Notropis volucellus	L. teres	0	Etheostoma radiosum	L. teres	0.625
Notropis volucellus	M. nervosa	0	Etheostoma radiosum	M. nervosa	0.375
Notropis volucellus	O. reflexa	0.375	Etheostoma radiosum	O. reflexa	0.875
Notropis volucellus	O. jacksoniana	0.375	Etheostoma radiosum	O. jacksoniana	0.875
Notropis volucellus	P. purpuratus	0.375	Etheostoma radiosum	P. purpuratus	0.875
Notropis volucellus	P. occidentalis	0.125	Etheostoma radiosum	P. occidentalis	0.75
Notropis volucellus	Q. pustulosa	0.375	Etheostoma radiosum	Q. pustulosa	0.875
Notropis volucellus	Q. quadrula	0.375	Etheostoma radiosum	Q. quadrula	0.875
Notropis volucellus	T. verrucosa	0.375	Etheostoma radiosum	T. verrucosa	0.875
Notropis volucellus	T. truncata	0.375	Etheostoma radiosum	T. truncata	0.875
Notropis volucellus	V. arkansasensis	0.125	Etheostoma radiosum	V. arkansasensis	-
Notropis volucellus	V. lienosa	0.25	Etheostoma radiosum	V. lienosa	0.25
Campostoma anomalum	A. ligamentina	0.875	Percina copelandi	A. ligamentina	0.75
Campostoma anomalum	A. plicata	0.875	Percina copelandi	A. plicata	0.75
Campostoma anomalum	A. wheeleri	0.625	Percina copelandi	A. wheeleri	0.625
Campostoma anomalum	E. lineolata	0.875	Percina copelandi	E. lineolata	0.75
Campostoma anomalum	F. flava	0.875	Percina copelandi	F. flava	0.75
Campostoma anomalum	L. radiata	0.75	Percina copelandi	L. radiata	0.625
Campostoma anomalum	L. ovata	0.875	Percina copelandi	L. ovata	0.75
Campostoma anomalum	L. teres	0.5	Percina copelandi	THE RESERVE TO SERVE	0.5
Campostoma anomalum	M. nervosa	0.25	Percina copelandi	L. teres M. nervosa	0.25
Campostoma anomalum	O. reflexa	0.875	Percina copelandi	O. reflexa	0.75
Campostoma anomalum	O. jacksoniana	0.875	Percina copelandi	O. jacksoniana	0.75
Campostoma anomalum	P. purpuratus	0.875	Percina copelandi	THE PROPERTY OF THE PROPERTY O	0.75
Campostoma anomalum	P. occidentalis	0.625	Percina copelandi	P. purpuratus P. occidentalis	0.7

Table 11. Results of Jaccard Index for Fish and Mussel Presence/Absence.

Campostoma anomalum	Q. pustulosa	0.875	Percina copelandi	Q. pustulosa	0.75
Campostoma anomalum	Q. quadrula	0.875	Percina copelandi	Q. quadrula	0.75
Campostoma anomalum	T. verrucosa	0.875	Percina copelandi	T. verrucosa	0.75
Campostoma anomalum	T. truncata	0.875	Percina copelandi	T. truncata	0.75
Campostoma anomalum	V. arkansasensis	0.25	Percina copelandi	V. arkansasensis	0.25
Campostoma anomalum	V. lienosa	0.375	Percina copelandi	V. lienosa	0.375
Pimephales notatus	A. ligamentina	0.75	Percina sciera	A. ligamentina	0.75
Pimephales notatus	A. plicata	0.75	Percina sciera	A. plicata	0.75
Pimephales notatus	A. wheeleri	0.5	Percina sciera	A. wheeleri	0.5
Pimephales notatus	E. lineolata	0.75	Percina sciera	E. lineolata	0.75
Pimephales notatus	F. flava	0.75	Percina sciera	F. flava	0.75
Pimephales notatus	L. radiata	0.625	Percina sciera	L. radiata	0.625
Pimephales notatus	L. ovata	0.75	Percina sciera	L. ovata	0.75
Pimephales notatus	L. teres	0.5	Percina sciera	L. teres	0.5
Pimephales notatus	M. nervosa	0.375	Percina sciera	M. nervosa	0.375
Pimephales notatus	O. reflexa	0.75	Percina sciera	O. reflexa	0.75
Pimephales notatus	O. jacksoniana	0.75	Percina sciera	O. jacksoniana	0.75
Pimephales notatus	P. purpuratus	0.75	Percina sciera	P. purpuratus	0.75
Pimephales notatus	P. occidentalis	0.625	Percina sciera	P. occidentalis	0.625
Pimephales notatus	Q. pustulosa	0.75	Percina sciera	Q. pustulosa	0.75
Pimephales notatus	Q. quadrula	0.75	Percina sciera	Q. quadrula	0.75
Pimephales notatus	T. verrucosa	0.75	Percina sciera	T. verrucosa	0.75
Pimephales notatus	T. truncata	0.75	Percina sciera	T. truncata	0.75
Pimephales notatus	V. arkansasensis	0.25	Percina sciera	V. arkansasensis	0.375
Pimephales notatus	V. lienosa	0.125	Percina sciera	V. lienosa	0.25

Class Value	#Pixels	%Total Pixels	Class Description
1	32274	0.33	Urban
2	16667	0.17	Primary Roads
3	2360400	24.05	Pasture/Regrowth/Cropland
4	1099017	11.20	Deciduous Forest
5	1438093	14.65	Coniferous Forest
6	4578552	46.65	Mixed Forest
7	41738	0.43	Rivers/Streams/Wetland
8	156394	1.59	Reservoirs
9	91128	0.93	Clouds/Cloud Shadows

Table 12. Number and percentage of pixels for each category in the final classification.

	L		Re	feren	ce Da	ita			100		
Classified Data	1	2	3	4	5	6	7	8	Totals	User's Accuracy	Producer's Accuracy
1. Urban	20	1	0	0	0	0	0	0	21	95.2%	100.0%
2. Primary Roads	0	0	0	0	0	0	0	0	0	******	0.0%
3. Pasture/Regrowth/Cropland	0	14	17	1	0	2	0	0	34	50.0%	85.0%
4. Deciduous Forest	0	1	3	3	0	2	0	0	9	33.3%	13.6%
5. Coniferous Forest	0	0	0	2	13	4	9	1	29	44.8%	35.0%
6. Mixed Forest	0	6	0	16	7	13	2	2	46	28.3%	61.9%
7. Rivers/Streams/Wetland	0	0	0	0	0	0	8	0	8	100.0%	34.8%
8. Reservoirs	0	0	0	0	0	0	4	17	21	81.0%	85.0%
Totals	20	22	20	22	20	21	23	20	168	L'ETTE.	

Table 13. Error matrix showing both user's and producer's accuracy estimates along with overall accuracy for all categories except for Clouds/Cloud Shadows.

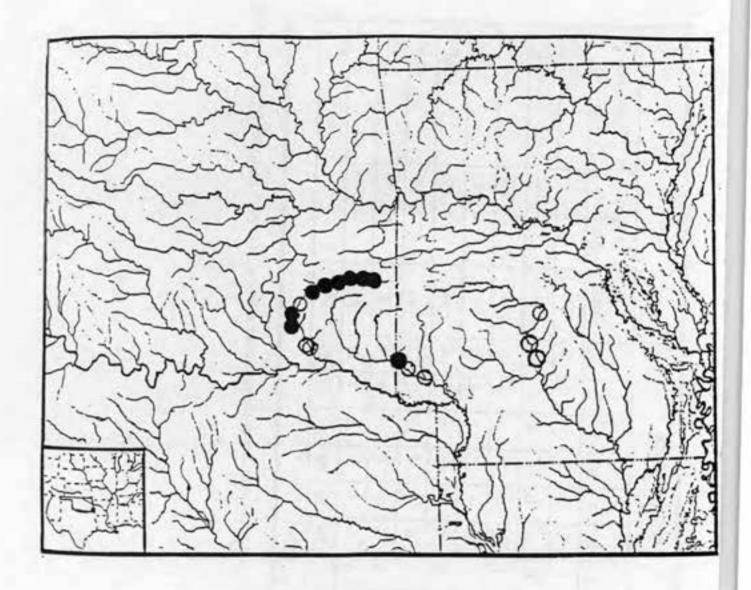
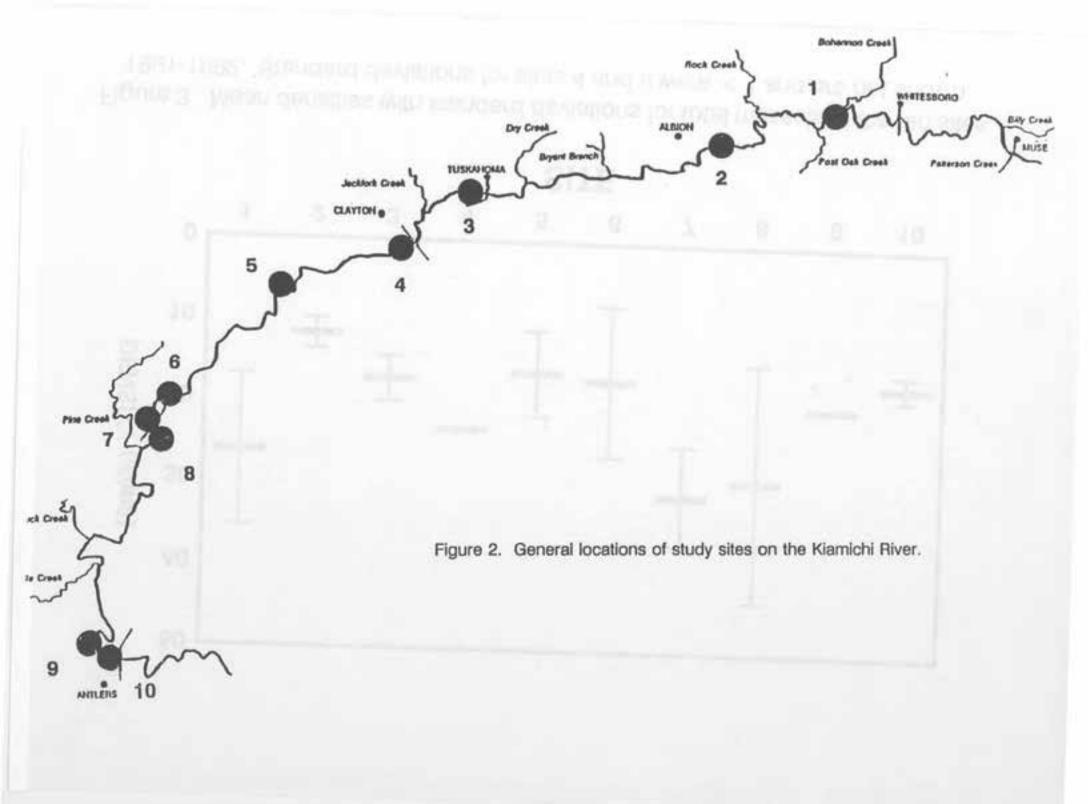


Figure 1. Historical and present distribution of Arkansia wheeleri. Solid circles indicate extant sites and open circles represent extirpated sites. In this figure sites in close proximity to one another appear as one dot.



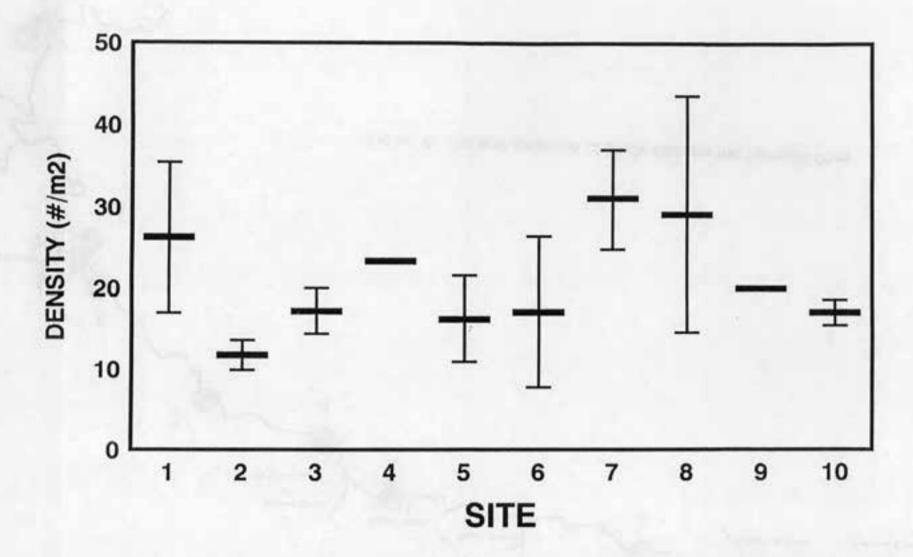


Figure 3. Mean densities with standard deviations for total mussels at the ten sites, 1991-1992. Standard deviations for sites 4 and 9 were < 1 and are not shown.

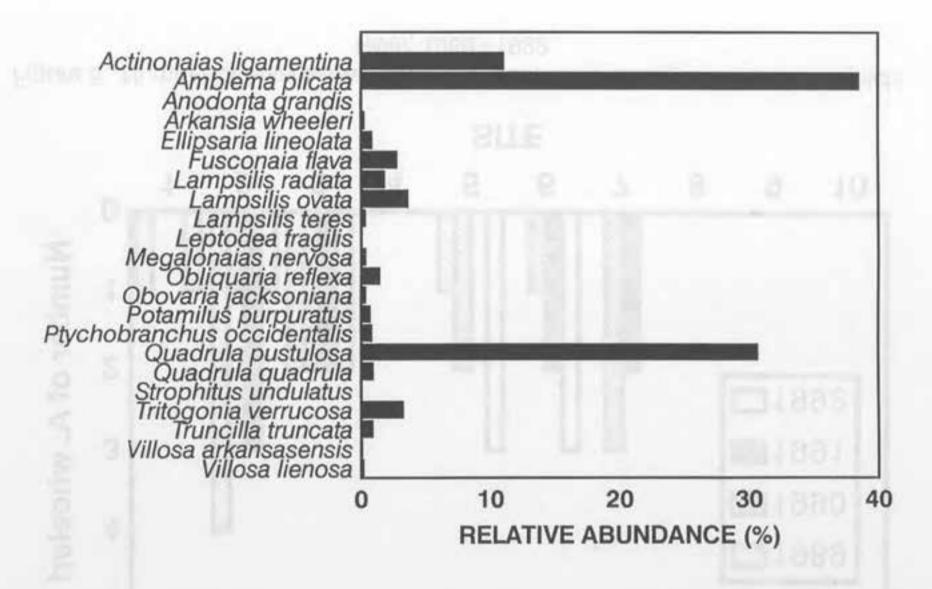


Figure 4. Mean relative abundance of mussel species in the Kiamichi River 1990-92.

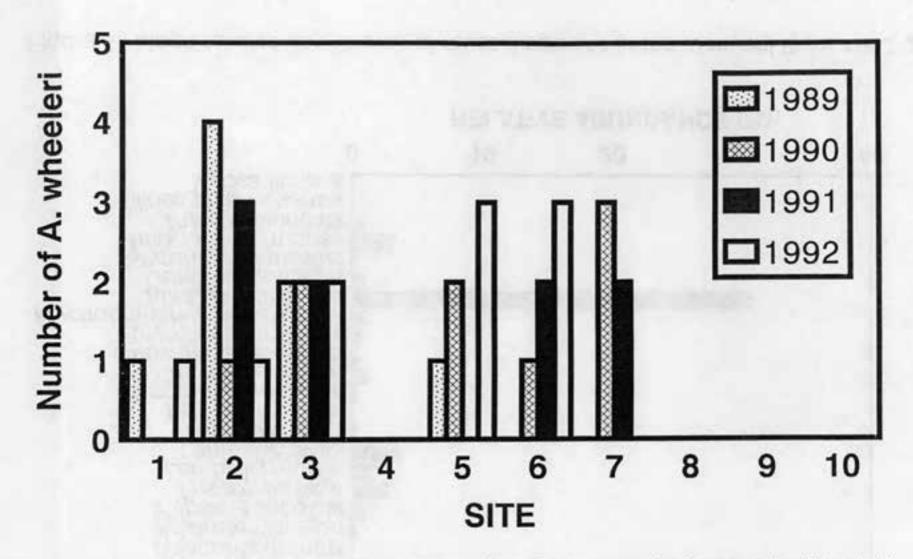


Figure 5. Number of Arkansia wheeleri found at the ten study sites on the Kiamichi River, 1989 - 1992.

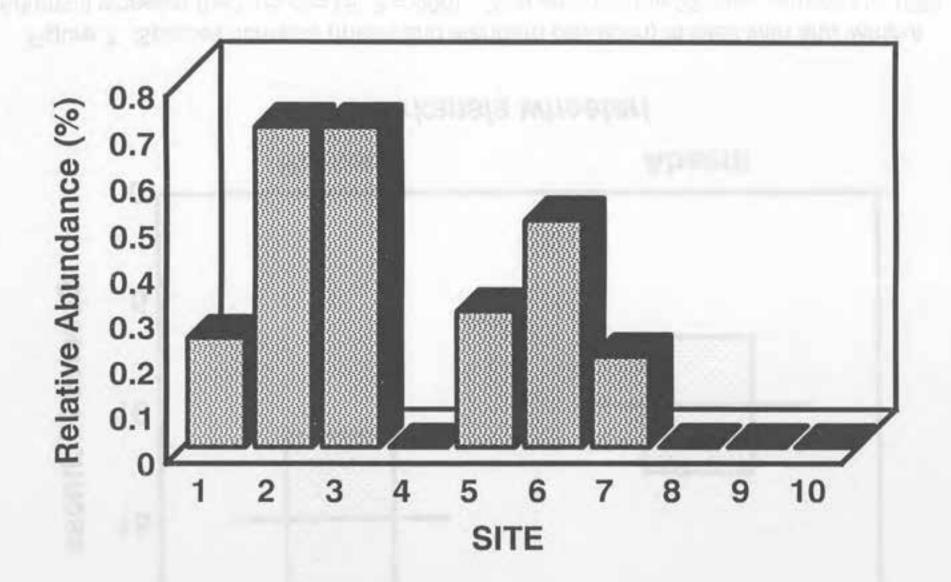


Figure 6. Relative abundance of Arkansia wheeleri at ten sites in the Kiamichi River, 1990-1992.

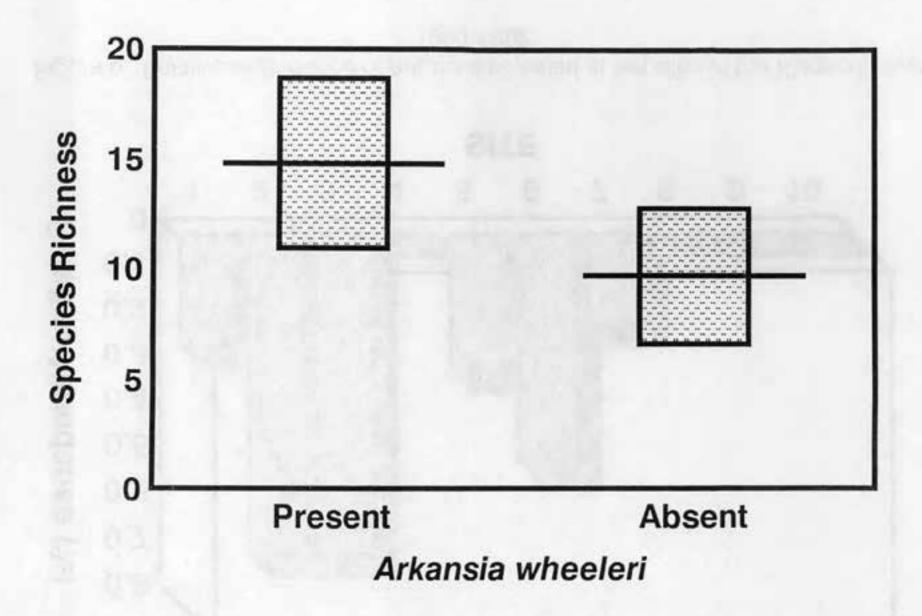
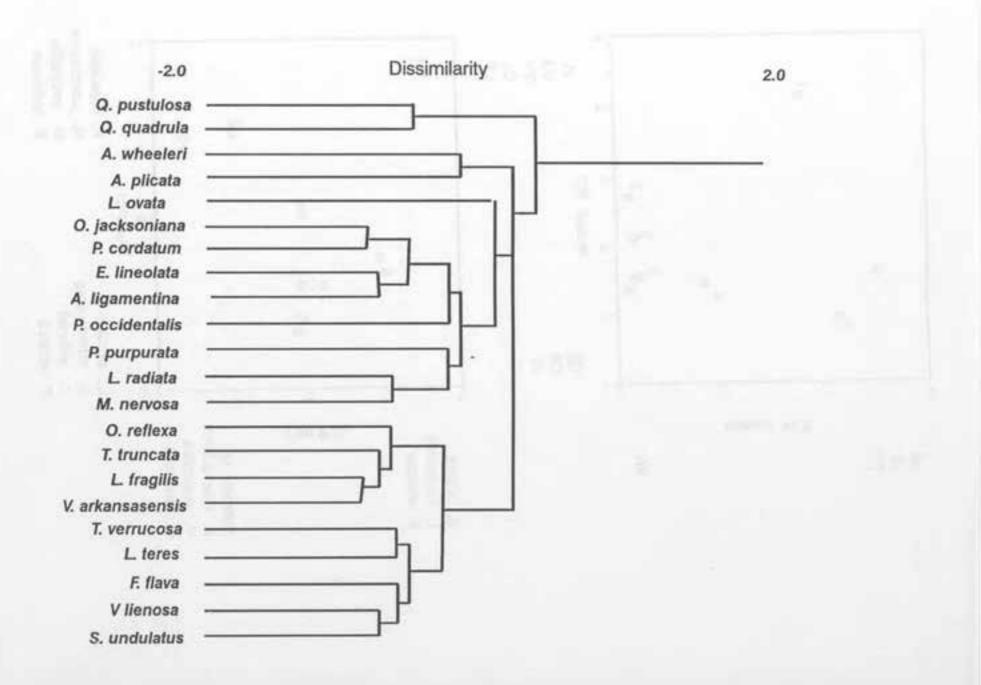


Figure 7. Species richness (mean and standard deviation) at sites with and without Arkansia wheeleri (t=3.18, df=15, P=.006). Data are from the 22 sites sampled in 1990.

Figure 8. Cluster analysis for 22 sites in the Kiamichi River sampled in 1990.



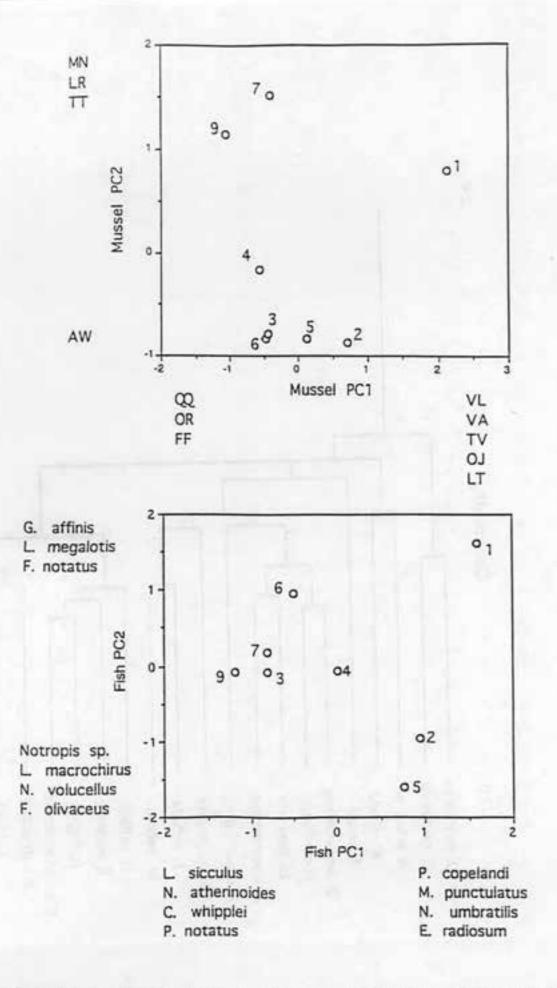


Figure 9. Principal components ordination. Data are means from 1991-1992 for eigsites.

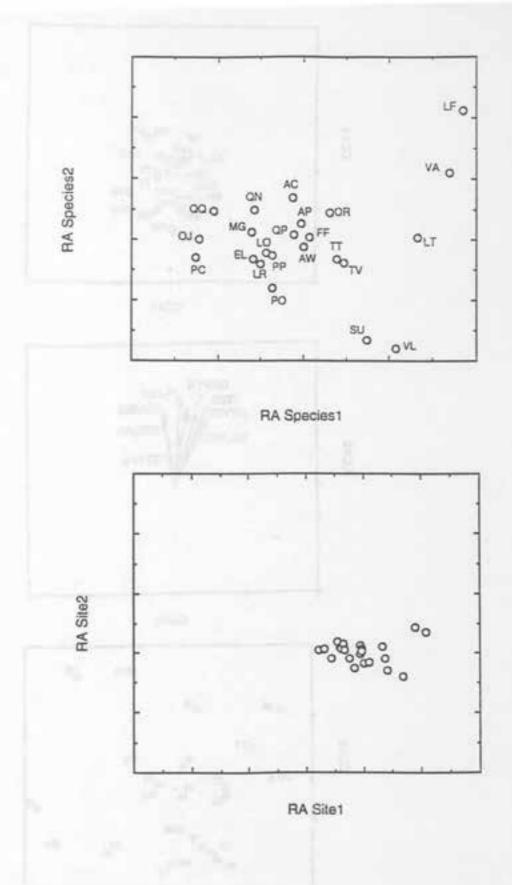


Figure 10. Reciprocal averaging ordination. Data are means from 1991-1992 for eight sites.

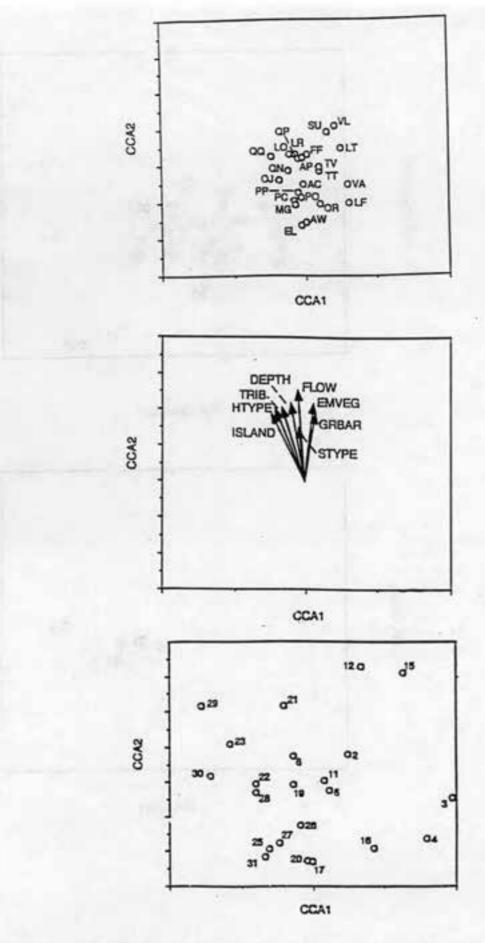


Figure 11. Results of canonical correspondence analysis for data from 1990. The top graph shows the approximate centers of species distributions along the first two CCA axes, the middle graph shows the location of habitat vectors along the axes, at the bottom graph shows the positions of the twenty two sites.

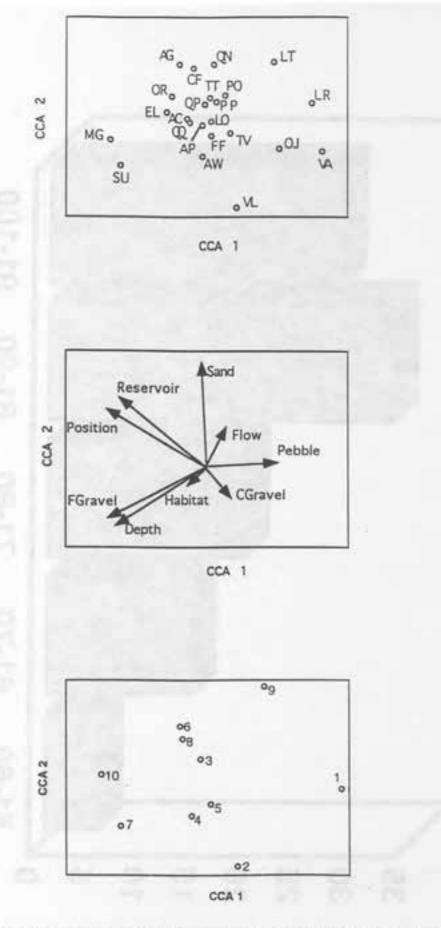


Figure 12. Results of canonical correspondence analysis for data from 1991-1992. The top graph shows the approximate centers of species distributions along the first two CCA axes, the middle graph shows the location of habitat vectors along the axes, and the bottom graph shows the positions of the ten study sites. Data used in this analysis were means for 1991-1992.

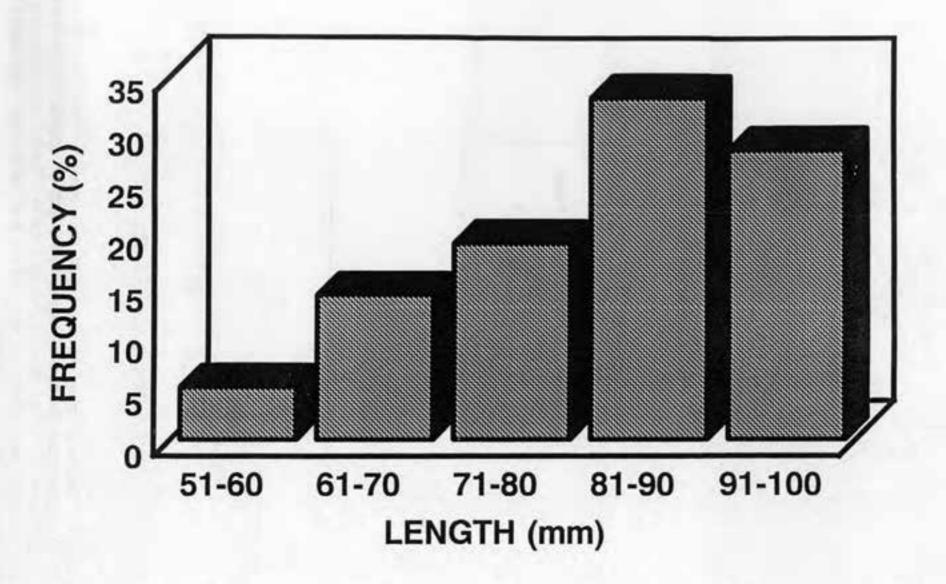


Figure 13. Total lengths of live Arkansia wheeleri from the Kiamichi River, 1989 - 1992.

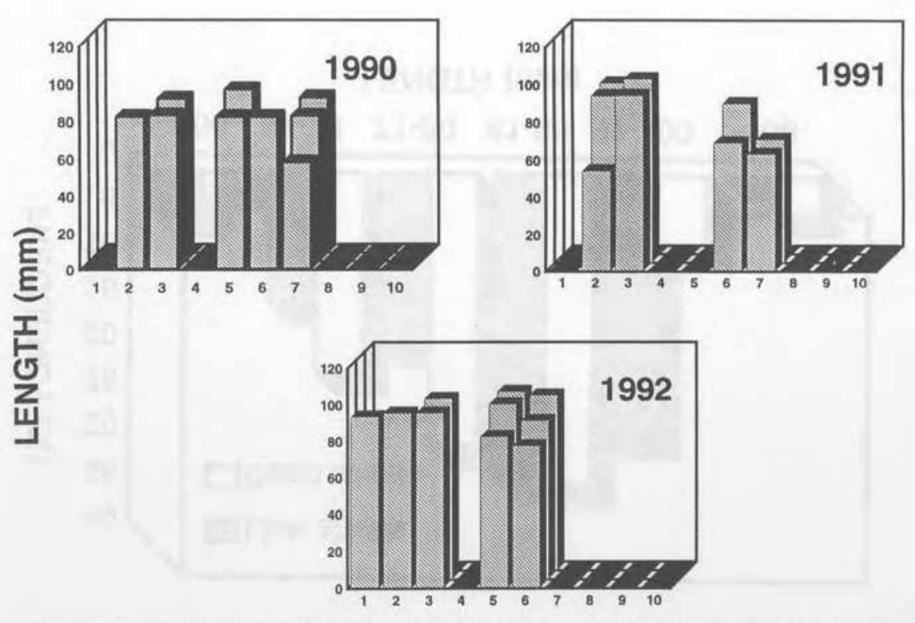


Figure 14. Lengths (mm) of Arkansia wheeleri found at ten study sites on the Kiamichi River 1990-92. Each bar represents an individual mussel.

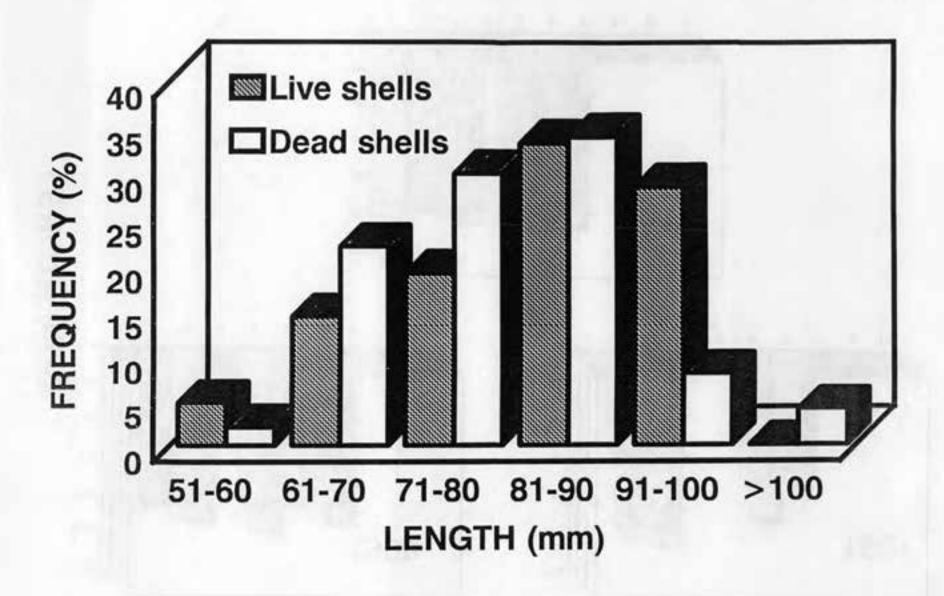


Figure 15. Total lengths of live Arkansia wheeleri compared to relict shells from the Kiamichi River (t=1.9, df=78, P=0.03).

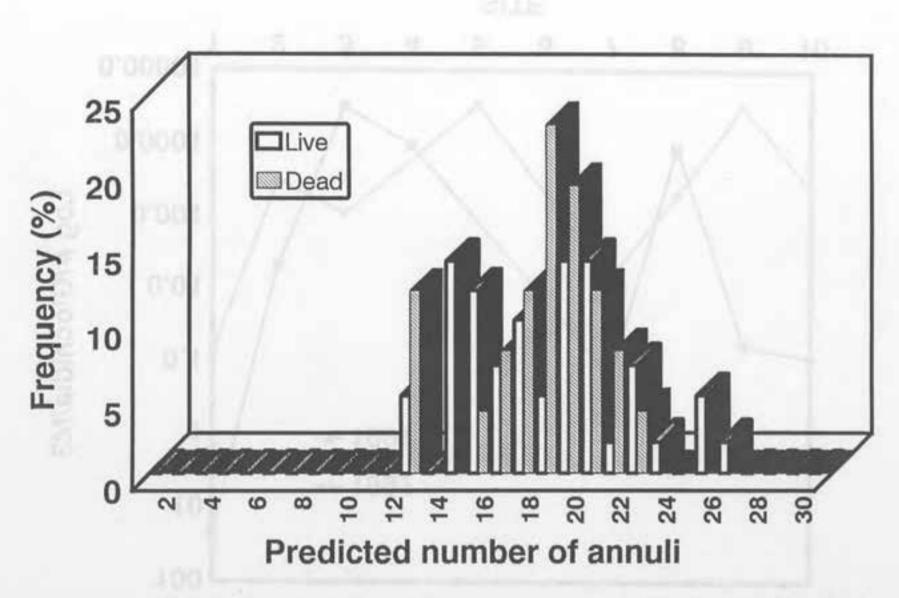


Figure 16. Predicted number of annuli for live Arkansia wheeleri versus relict shells from the Kiamichi River (t=0.84, df=54, P=0.19).

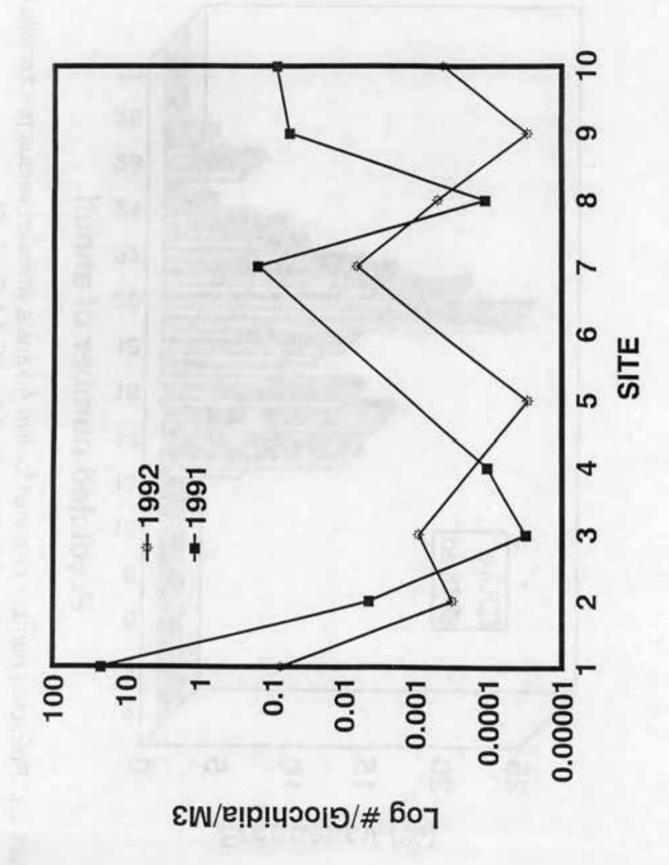


Figure 17. Densities of drifting glochidia.

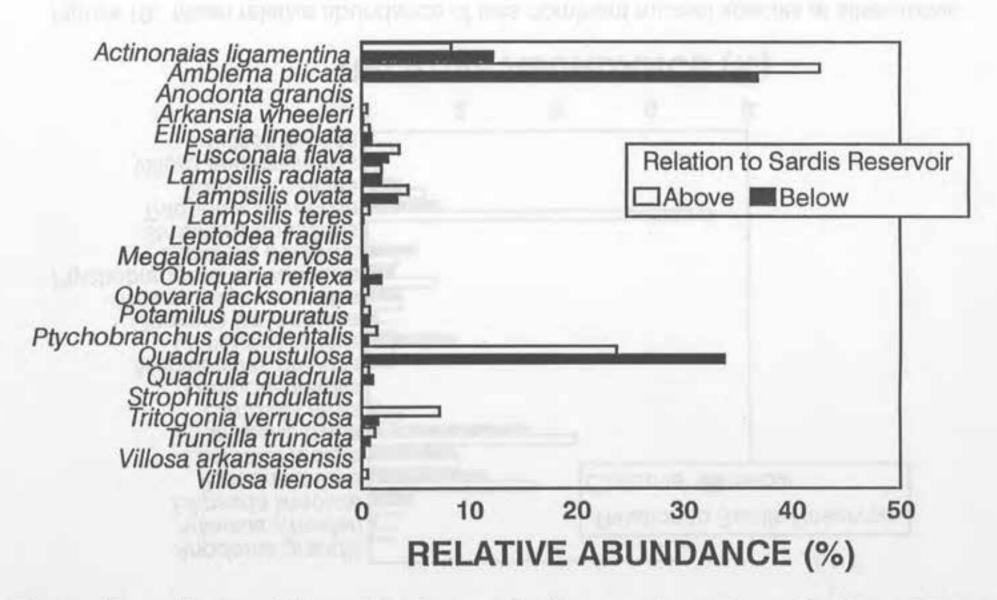


Figure 18. Mean relative abundance of all mussel species at sites above and below Sardis Reservoir, 1990-1992.

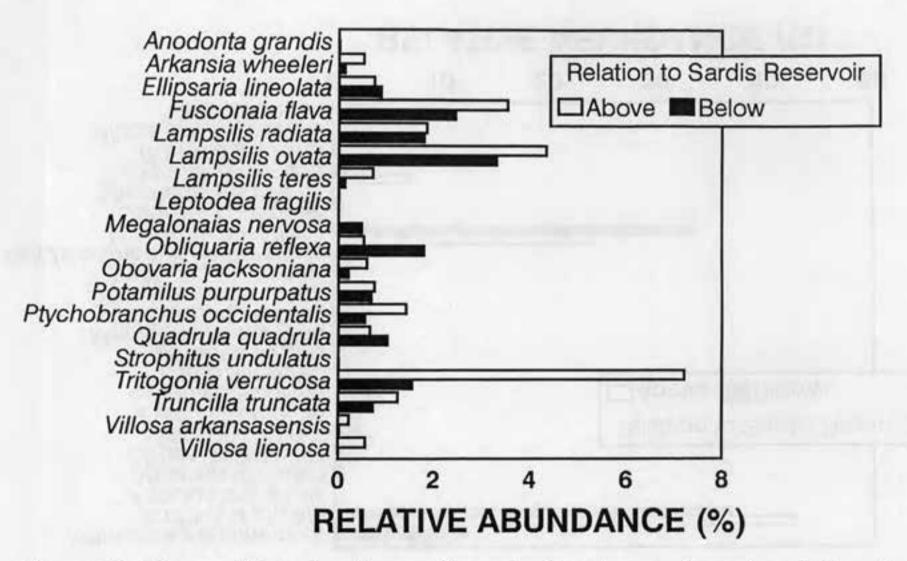


Figure 19. Mean relative abundance of less dominant mussel species at sites above and below Sardis Reservoir, 1990-92.

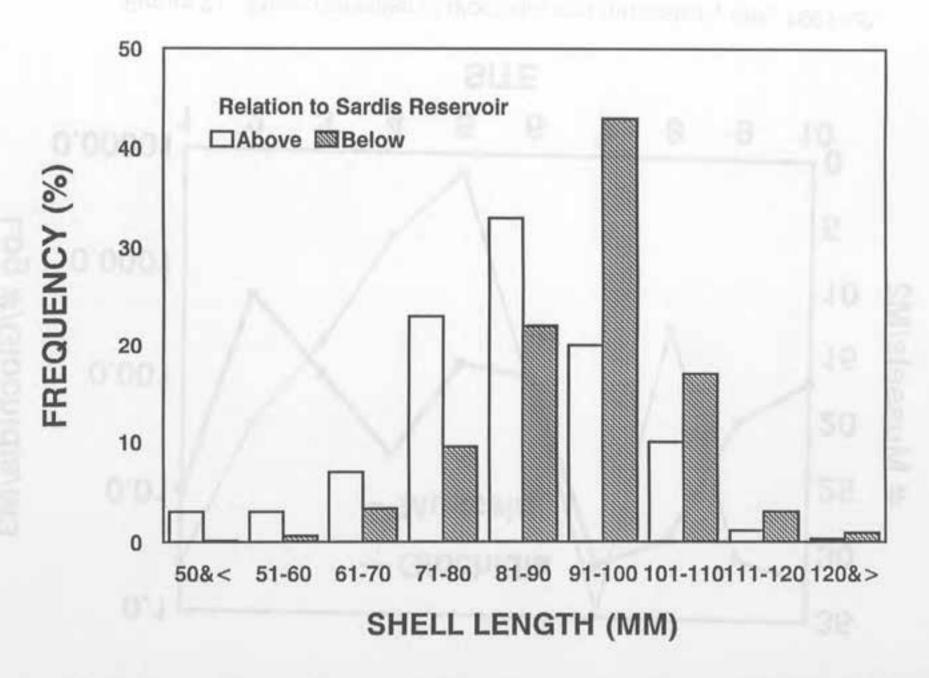


Figure 20. Lengths (mm) of live Amblema plicata from sites above and below Sardis Reservoir in 1991.

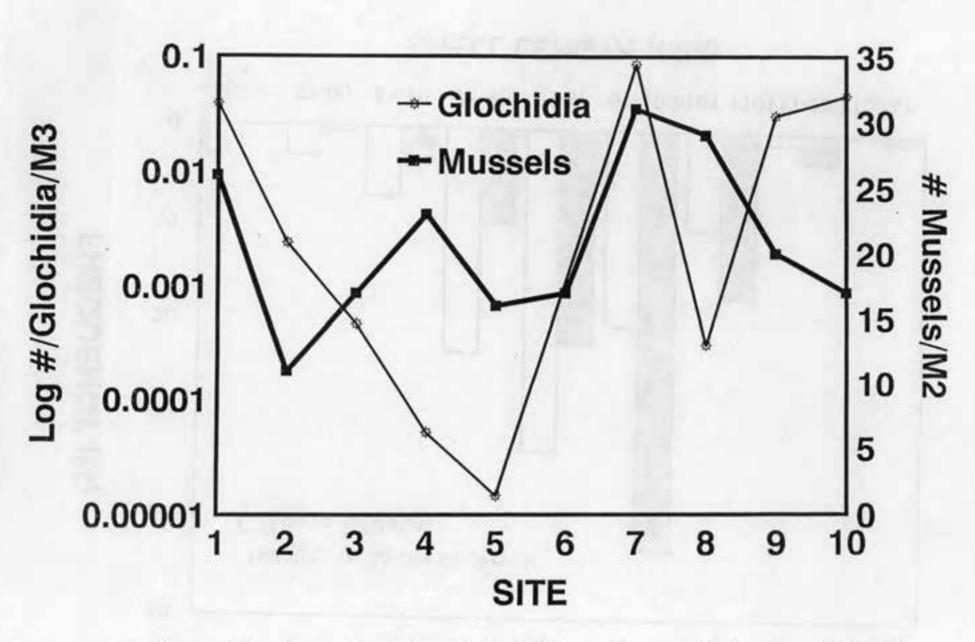
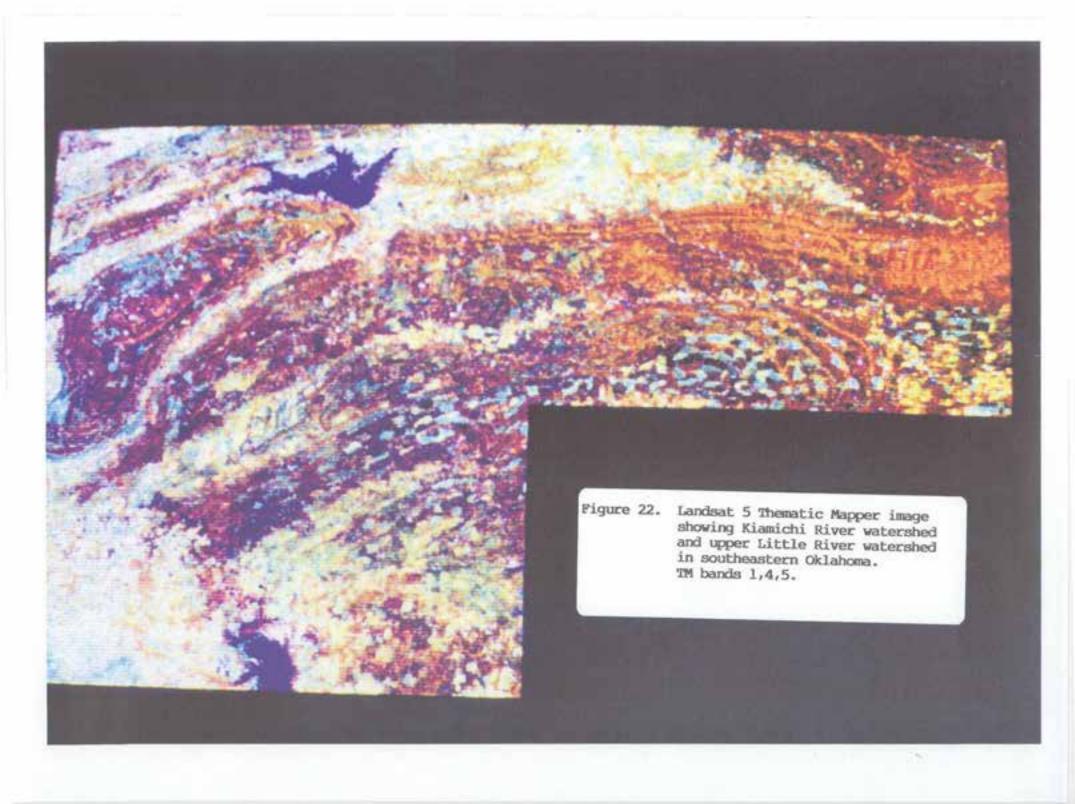
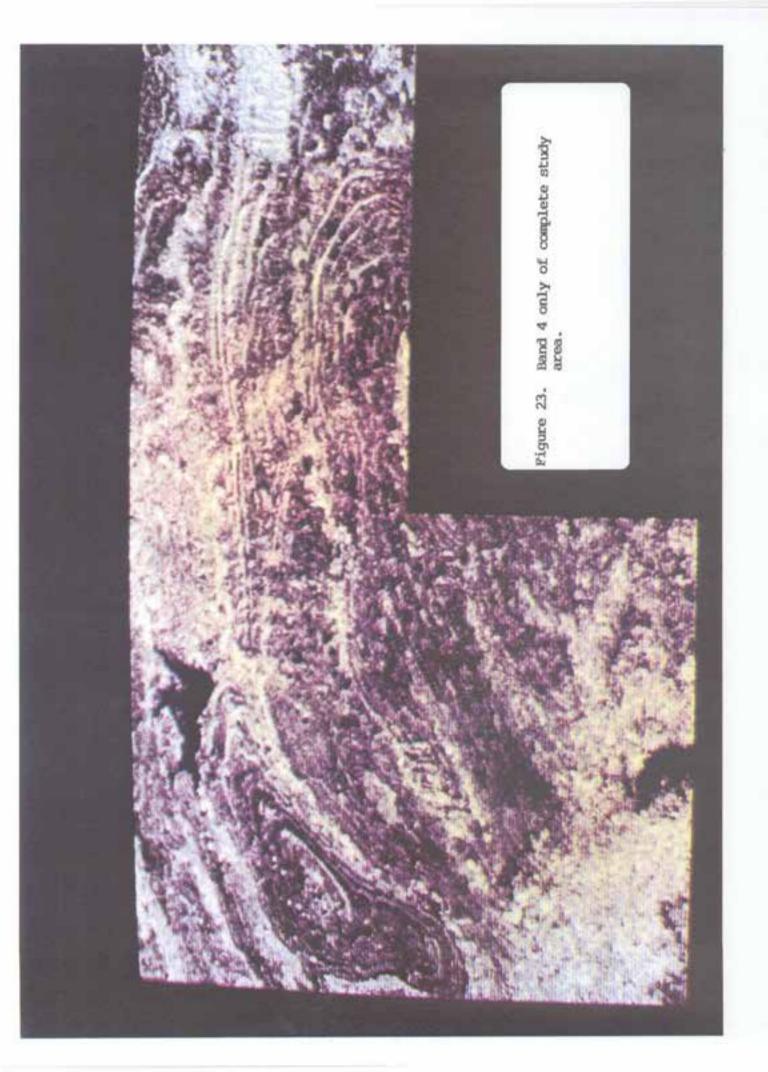


Figure 21. Mean densities of glochidia and mussels by site, 1991-92.





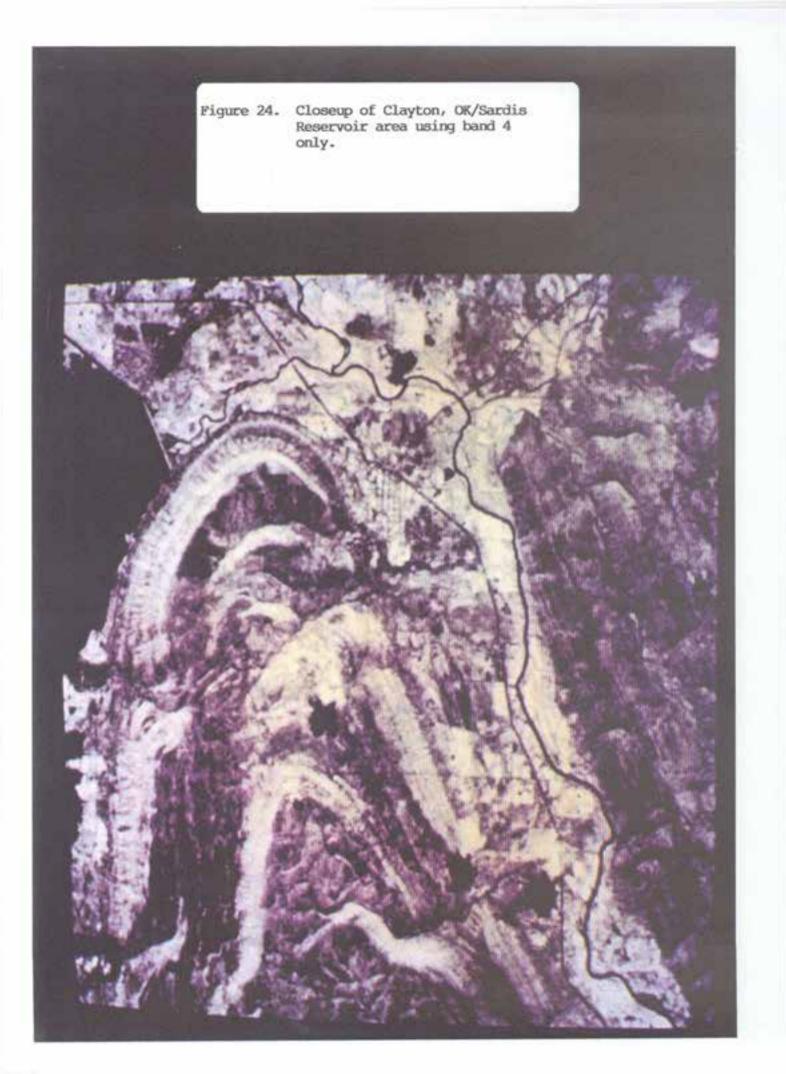


Figure 25. Map colors and categories associated with final classi-fication.

Urban Primary Roads Pasture/Regrowth/Cropland Deciduous Forest Coniferous Forest Mixed Forest Rivers/Streams/Wetlands Reservoirs

Clouds/Cloud Shadows

Landuse Classification for Kiamichi River Watershed, OK Urban Primary Roads Pasture/Regrowth/Cropland Deciduous Forest Coniferous Forest

Figure 26. Final classification of Kiamichi River and upper Little River watersheds.



APPENDIX 1

ADDITIONAL WATER QUALITY DATA

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07335700 KIAWICHI RIVER NEAR BIG CEDAR, OK (Hydrologic benchmark station)

LOCATION -- Lat 34038'18*, long 94036'45*, in SW 1/4 SE 1/4 sec.18, T.2 N., R.25 E., Le Flore County, Hydrologic Unit 11148185, Suachita National Forest, on downstream side of right bank pier of bridge on State Highway 63, 8.2 si upstream from Rattlesh Creek, 1.1 si upstream from Big Branch, 2.1 mi east of Big Cedar, and at mile 157.6.

DRAINAGE AREA .-- 48.1 m/2

WATER-DISCHARGE RECORDS

PERIOD OF RECORD -- October 1965 to current year

GAGE .-- Water-stage recorder Datum of gage is 886.97 ft above National Geodetic Vertical Datum of 1929

REMARKS -- Records good

UNPUBLISHED RECORDS

AVERAGE DISCHARGE -- 26 years, 82 7 ft3/s, 28 8 in/yr, 59,928 acre-ft/yr

EXTREMES FOR PERIOD OF RECORD. -- Maximum discharge, 27,488 ft³/s, May 19, 1998, gage height, 19.68 ft; from rating curve exter above 9,888 ft³/s; no flow at times in most years.

EXTREMES FOR CURRENT YEAR .-- Peak discharges greater than base discharge of 2,000 ft2/s and maximum (+):

Date	Tine	Discharge (ft /s)	Gage Height (ft)	Date	Tine	Oischarge (ft ^a /s)	Gage Height (ft)
Oct. 7	1988	2,278	9 52	Apr. 13	1645	18,188	14.51
Oct. 8	1588	8,918	13 99	Way 3	8415	2,188	9.85
War. 22	8815	•11,288	•14 98	July 27	1515	3,938	18.98
Apr. 12	8638	8,128	13 62	July 28	8738	18,288	14.56

Winisum daily discharge, 8 68 ft3/s July 28.

DISCHARGE, CUBIC FEET PER SECOND, WATER YEAR OCTOBER 1998 TO SEPTEMBER 1991 DAILY MEAN VALUES

DAY	OCT	NOV	DEC	JAN	FEB	WAR	APR	MAY	JUN	JUL	AUG	SEP
1 2 3 4 5	7.8 5.5 8.8 28 21	9.8 11 11 57 82	67 79 227 147 114	197 188 159 133 128	33 38 29 28 41	154 146 184 84 72	124 188 124 183 88	83 87 882 297 258	28 22 18 13 11	7.2 5.9 5.9 8.9 6.1	58 35 26 19 14	2.8 2.3 2.6 2.4 6.5
6 7 8 9 18	17 577 1848 1488 325	58 51 45 141 124	92 75 63 53 42	533 413 217 177 486	96 85 88 73 65	64 54 46 48 35	82 159 185 148 169	178 122 95 76 64	8.2 7.5 6.4 5.7	4 9 4 2 3.6 3.1 2.6	18 8 3 5.5 5.5 4 4	5.9 6.1 28 17 12
11 12 13 14 15	185 185 75 66 43	97 77 64 53 46	39 37 33 38 28	314 212 163 132 398	57 52 51 45 38	32 31 26 26 24	98 1888 3248 989 483	53 43 36 38 28	16 12 9.1 7.5 7.7	2.3 1.9 1.6 1.5 1.3	3.9 3.3 4.5 5.4 3.4	9.8 7.3 5.2 5.4 4.7
16 17 18 19 28	35 38 25 21 18	39 34 31 28 26	29 13 65 99	383 288 172 172 145	35 35 34 31 28	22 44 39 35 35	246 193 958 325 285	27 24 19 19 24	56 34 22 17 13	1.2 1.1 1.8 97 89	3.4 4.5 4.8 2.9 2.4	5.8 22 29 28 24
21 22 23 24 25	25 23 19 16 14	28 27 24 22 28	112 101 86 77 66	122 188 95 81 78	26 24 22 21 21	2378 459 237 161	153 142 168 83 84	41 45 58 43 168	9.9 73 159 55 37	81 .78 .68 164 132	2.5 2.7 2.1 1.8 1.5	28 16 13 11 9.3
26 27 28 29 38 31	13 11 11 9.8 9.3 9.1	23 63 99 85 75	68 77 94 652 627 281	62 56 52 45 42 37	19 18 17	126 215 174 249 284 157	68 157 153 149 187	182 78 51 39 31 25	28 21 16 13 9.5	61 775 2388 286 122 73	1.5 1.5 2.1 1.6 1.6 2.3	7.6 6.5 5.7 5.2 4.6
TOTAL MEAN MAX WIN AC-FT CFSM IN.	5835.5 162 1848 6.5 9998 4.85 4.67	1558.8 51.7 141 9.8 3888 1.29 1.44	3655 118 652 13 7258 2.94 3.39	5532 178 533 37 18978 4 45 5.13	1135 40.5 96 17 2250 1.01 1.05	5517 178 2378 22 18948 4 44 5 12	18859 362 3248 68 21548 9 83 18.67	3898 99.7 882 19 6138 2.49 2.87	737.5 24.6 159 5.7 1468 .61	3981.43 128 2388 .68 7988 3.28 3.69	237.6 7.66 58 1.5 471 .19	318.1 18.6 29 2.3 631 26 .38
	1 2 3 4 5 6 7 8 9 8 11 12 3 14 15 16 7 18 19 28 21 22 23 24 5 10 TALL WALL OF SM WALL OF	1 7.8 2 5.5 3 8.8 4 28 5 21 6 17 7 577 8 1648 9 1488 18 325 11 185 12 185 13 75 14 56 15 43 16 35 17 38 18 25 19 21 28 18 21 22 22 23 23 19 24 16 25 14 26 13 27 11 28 18 21 26 22 23 23 19 24 16 25 14 26 13 27 11 28 18 27 28 18 27 28 18 28 29 38 31 9.3 31 9.1 TOTAL \$836.5 UEAN 1648 UIN 6.5 6.5 UEAN 1648 UIN 6.5 UEAN 1648 UIN 6.5 UIN 6.	1 7 8 9 8 2 6 5 11 3 8 8 11 4 28 57 5 21 82 6 17 68 7 577 51 8 1848 45 9 1488 141 18 325 124 11 185 97 12 185 77 13 75 64 14 14 56 53 15 43 46 16 35 39 17 38 34 18 25 31 19 21 28 28 18 26 21 25 28 22 23 27 23 19 24 24 16 22 25 14 28 26 13 23 27 11 63 28 19 24 24 16 22 25 14 28 26 13 23 27 11 63 28 19 24 24 16 22 25 14 28 26 13 23 27 11 63 28 19 24 24 16 22 25 14 28 26 13 23 27 11 63 28 19 24 24 16 22 25 14 28 26 13 23 27 11 63 28 11 99 29 9 8 85 38 9 3 75 31 9 1 TOTAL 5836 5 1558 8 UEAN 1848 141 UIN 6 5 9 8 AC-FT 9998 3888 CFSW 4 85 1.29	1 7.8 9.8 67 2 6.5 11 79 3 8.8 11 227 4 28 57 147 5 21 62 114 6 17 68 92 7 577 51 75 8 1848 45 63 9 1468 141 53 18 325 124 42 11 185 97 39 12 185 77 37 13 75 64 33 14 56 53 38 15 43 46 28 16 35 39 29 17 38 34 13 18 25 31 65 19 21 28 99 28 18 26 98 21 25 28 112 22 23 27 181 28 19 24 86 24 16 22 77 25 14 28 68 26 13 23 68 27 11 63 77 28 11 99 94 29 9.8 85 652 38 9.3 75 627 31 9.1 281 TOTAL 5836.5 1558.8 3655 UEAN 1648 141 652 UIN 6.5 9.8 13 AC-FT 9990 3888 7258 CFSW 4.85 1.29 2.94	1 7.8 9.8 67 197 2 6 5 11 79 188 3 8 8 11 227 159 4 28 57 147 133 5 21 62 114 128 6 17 68 92 533 7 577 51 75 413 8 1848 45 63 217 9 1468 141 53 177 18 325 124 42 466 11 185 97 39 314 12 185 77 37 212 13 75 64 33 163 14 56 53 38 132 15 43 46 28 398 16 35 39 29 383 17 38 34 13 288 18 25 31 55 172 19 21 28 99 172 28 18 26 98 148 21 25 28 112 122 22 23 27 161 168 21 26 28 112 122 22 23 27 161 168 23 19 24 86 95 24 16 22 77 51 25 14 28 66 78 26 13 23 68 82 27 11 63 77 56 28 11 99 94 55 29 9.8 85 652 45 38 9.3 75 627 42 31 9.1 281 37 TOTAL 5836.5 1558.8 3655 5532 UEAN 162 51.7 118 178 UAX 1848 141 652 533 UIN 165 98 13 37 AC-FT 9990 3888 7258 18978 CFSW 4.85 1.29 2.94 4 45	1 7.8 9.8 67 197 33 2 5.5 11 79 188 38 3 8.8 11 227 159 29 4 28 57 147 133 28 5 21 62 114 126 41 6 17 68 92 533 96 7 577 51 75 413 86 8 1648 45 63 217 88 9 1488 141 53 177 73 18 325 124 42 486 65 11 185 97 39 314 57 12 185 77 37 212 52 13 75 64 33 163 51 14 15 53 39 29 383 35 15 43 46 28 398 38 16 35 39 29 383 35 17 38 34 13 288 35 18 25 31 55 172 34 18 25 31 55 172 34 19 21 28 99 172 34 19 21 28 99 172 34 28 18 26 98 145 28 21 25 26 112 122 26 22 23 19 24 86 95 22 24 16 22 77 81 21 25 14 28 68 78 21 26 13 23 68 82 19 27 11 63 77 56 18 28 11 99 94 52 17 29 18 56 57 42 21 25 26 18 77 56 18 26 13 23 68 82 19 27 11 63 77 56 18 28 11 99 94 52 17 29 18 56 52 45 31 9.1 281 37 TOTAL 5835.5 1558.8 3655 5532 1135 UEAN 162 51.7 118 178 48.5 UAX 1846 141 652 533 96 UIN 6.5 9.8 13 37 TOTAL 5835.5 1558.8 3655 5532 1135 UEAN 162 51.7 118 178 48.5 UAX 1846 141 652 533 96 UIN 6.5 9.8 13 37 TOTAL 5835.5 1558.8 3655 5532 1135 UEAN 162 51.7 118 178 48.5 UAX 1846 141 652 533 96 UIN 6.5 9.8 13 37 77 AC-FT 9996 3888 7258 18978 2258 CFSW 4.85 1.29 2.94 4.45 1.81	1 7.8 9.8 67 197 33 154 2 6 5 11 79 188 38 146 3 8.8 11 227 159 29 184 4 28 57 147 133 28 64 5 21 62 114 128 41 72 6 17 68 92 533 96 64 7 577 51 75 413 85 54 8 1848 45 63 217 88 46 9 1488 141 53 177 73 48 18 325 124 42 486 65 35 11 185 97 39 314 57 32 12 185 77 37 212 52 31 13 75 64 33 163 51 28 14 55 63 38 132 45 26 15 43 46 28 398 38 24 16 35 39 29 383 35 22 17 38 34 13 288 35 44 18 25 31 65 77 37 212 52 31 18 25 31 55 77 37 212 52 31 19 24 86 28 398 38 24 16 35 39 29 383 35 22 17 38 34 13 288 35 44 18 25 31 65 77 37 31 35 28 18 25 98 145 28 35 21 26 28 112 122 26 489 24 16 22 77 81 21 237 25 14 28 66 78 21 161 26 13 23 68 52 19 126 27 11 63 77 55 18 21 237 28 19 24 86 95 22 489 24 16 22 77 81 21 237 25 14 28 66 78 21 161 26 13 23 68 52 19 126 27 11 63 77 55 18 215 28 11 99 94 52 17 174 29 9.8 85 652 45 249 38 9.3 75 627 42	1 7.8 9.8 67 197 33 154 124 2 6.5 11 79 188 38 146 188 3 8.8 11 227 159 29 184 124 4 28 57 147 133 28 84 183 5 21 82 114 128 86 54 159 8 1848 45 63 217 88 46 185 9 1488 141 53 177 73 48 148 18 325 124 42 426 65 35 169 11 185 97 39 314 57 32 98 12 185 77 37 212 52 31 1888 13 75 64 33 163 51 26 3248 14 56 53 38 132 45 26 989 15 43 46 28 398 38 24 483 16 35 39 29 383 35 22 246 17 38 34 13 288 35 44 193 18 25 124 86 69 172 34 39 958 18 25 31 65 172 34 39 958 19 21 28 99 172 31 35 325 28 18 26 98 145 28 35 225 21 26 28 112 122 26 48 153 22 23 27 181 188 24 2378 142 23 19 24 86 95 172 34 39 958 18 25 31 65 172 34 39 958 18 26 98 145 28 35 225 21 26 28 112 122 26 48 153 22 23 27 181 188 24 2378 142 23 19 24 86 95 22 469 188 24 16 22 77 81 21 237 83 25 14 28 66 95 22 469 188 26 13 23 68 82 19 126 68 27 11 63 77 56 18 215 157 28 11 99 94 52 17 174 153 29 9 8 85 657 42 244 147 31 9.1 281 37 157 TDTAL 5836.5 1558.8 3655 5532 1135 5517 18859 MEAN 162 51.7 118 178 48 5 178 362 MIN 6.5 9.8 13 37 72 68 MC-FT 9990 3888 7258 18978 2258 18948 2154 49 83 MIN 6.5 9.8 13 37 72 68 MC-FT 9990 3888 7258 18978 2258 18948 2154 49 83	1 7 8 9.8 67 197 33 154 124 83 25 55 11 79 188 38 146 108 67 3 8.8 11 227 159 29 104 124 882 4 28 57 147 133 25 84 183 297 5 21 82 114 128 41 72 88 258 6 17 68 92 533 96 64 82 178 7 577 51 75 413 85 54 159 122 8 1848 45 63 217 88 46 185 95 1488 141 53 177 73 48 148 148 76 18 325 124 42 426 65 35 129 64 11 185 97 37 37 212 52 31 1888 43 177 18 18 325 124 42 426 65 35 129 64 11 185 97 37 31 212 52 31 1888 43 13 13 75 64 33 163 51 26 3248 35 14 14 15 15 17 18 17 18 18 18 18 18 18 18 18 18 18 18 18 18	1 7.8 9.8 67 197 33 154 124 83 28 2 5.5 11 79 188 38 146 188 67 22 3 8.8 11 227 159 29 184 124 882 18 4 28 57 147 133 28 84 183 297 13 5 21 82 114 128 41 72 88 258 11 6 17 68 92 533 98 64 82 178 18 7 577 51 75 413 86 54 159 122 8.2 8 1848 45 63 217 88 46 185 95 7.5 9 1488 141 53 177 73 48 148 148 75 6.4 18 325 124 42 486 65 35 169 64 5.7 11 185 97 39 314 57 32 98 53 1888 43 12 13 75 64 33 163 51 26 3248 36 9.1 14 56 53 38 132 45 26 3248 36 9.7 15 43 46 28 398 38 24 483 28 7.7 16 35 39 29 383 35 22 246 27 86 17 38 34 13 288 35 44 193 28 7.5 18 25 31 65 77 37 212 52 31 1888 43 12 13 75 64 33 163 51 26 3248 36 9.1 14 56 53 38 132 45 26 989 38 7.5 15 43 46 28 398 38 24 483 28 7.7 16 35 39 29 383 35 22 246 27 86 17 38 34 13 288 35 44 193 28 7.5 18 25 31 65 77 37 212 52 31 1888 35 9.1 19 21 28 99 172 31 35 325 19 17 28 18 25 31 65 77 37 18 18 18 35 325 19 17 29 18 26 28 112 122 28 48 183 35 325 19 17 20 18 26 28 112 122 28 48 183 35 325 19 17 21 26 28 112 122 28 48 183 35 325 19 17 22 23 27 181 188 24 2278 142 45 73 23 19 24 86 95 22 469 188 58 159 24 16 22 77 81 21 237 83 48 99 13 25 14 28 86 78 21 161 84 188 37 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 5836.5 1558.8 3655 5532 1135 5517 18859 3898 737.5 10TAL 58	1 7.8 9.8 67 197 333 154 124 83 28 7.2 2 5.5 11 79 188 38 146 188 67 22 5.9 3 8.8 11 227 159 29 184 124 882 18 5.9 4 28 57 147 133 26 84 183 297 13 8.9 5 21 82 114 128 41 72 88 258 11 6.1 6 17 68 92 533 96 64 82 178 188 24 237 8 1 1 6.1 6 17 68 92 138 297 13 8.9 7 577 51 75 413 86 54 159 122 8.2 4.2 8 1848 45 63 217 88 46 185 95 7.5 3.6 9 1408 141 53 177 73 48 148 75 6.4 3.1 18 325 124 42 486 65 35 169 64 5.7 2.6 11 185 97 39 314 57 32 98 53 16 2.3 12 185 77 37 212 52 31 1888 43 12 1.9 13 75 64 33 153 51 28 3248 36 9.1 1.6 14 56 53 38 132 45 26 28 3949 38 7.5 15 15 43 46 28 398 38 24 483 28 7.7 13 16 35 39 29 383 35 44 193 24 34 1.1 18 25 31 55 172 34 39 956 19 22 1.8 19 21 28 99 172 31 35 325 14 13 89 21 28 28 11 1 10 1 10 10 10 10 10 10 10 10 10 10	1 7.8 9.8 67 197 33 154 124 83 28 7.2 58 2 8.5 11 79 188 38 146 188 67 22 5.9 55 3 8.8 11 227 159 29 184 124 882 18 5.9 26 4 28 57 147 133 28 64 123 297 13 8.9 19 5 21 62 114 128 41 72 88 258 11 6.1 14 6 17 68 92 533 96 64 82 178 18 4 9 18 7 577 577 51 413 88 64 185 95 7.5 3.6 5.5 9 1468 445 63 217 88 46 185 95 7.5 3.6 5.5 9 1468 141 53 177 73 48 46 185 95 7.5 3.6 5.5 18 325 124 42 486 65 35 189 64 5.7 2.6 4 4 11 185 97 39 314 57 32 98 53 16 2.3 3.9 12 185 97 39 314 57 32 98 53 16 2.3 3.9 12 185 77 37 212 52 31 1888 43 12 19 3.3 13 75 64 33 163 51 26 3248 36 9.1 1.6 4.5 14 56 53 38 132 45 26 99 38 7.5 1.5 5.4 15 43 46 28 398 38 24 483 28 7.7 1.3 3.4 16 35 39 29 383 35 24 483 28 7.7 1.3 3.4 16 35 39 29 383 35 22 246 27 56 1.2 3.4 17 38 34 13 288 35 24 483 28 7.7 1.3 3.4 18 25 31 18 26 99 172 31 35 225 19 77 1.3 3.4 18 25 31 18 28 398 38 24 483 28 7.7 1.3 3.4 19 21 22 22 22 27 181 188 24 493 28 7.7 1.3 3.4 16 35 39 29 38 35 22 246 27 56 1.2 3.4 21 26 28 39 145 28 39 19 32 4 45 2 5 99 38 7.5 1.5 5.4 18 25 31 65 77 37 212 52 34 39 958 19 22 1.8 4.9 28 18 26 99 172 31 35 225 19 77 1.3 3.4 19 21 28 99 172 31 35 255 19 77 2.9 28 18 26 28 112 188 24 2378 142 45 73 78 22 1.8 4.9 29 21 28 99 172 31 35 255 19 77 1.3 8.9 24 18 25 31 165 172 34 39 958 19 22 1.8 4.9 28 18 26 99 172 31 35 255 19 17 77 2.9 28 18 26 28 112 188 24 2378 142 45 73 78 22 79 28 18 26 58 11 188 26 49 145 28 35 285 144 13 89 2.4 21 26 28 77 81 21 237 83 43 55 184 1.8 25 14 28 68 77 81 21 237 83 43 55 184 1.8 26 13 23 68 82 19 126 68 182 28 61 1.5 27 11 63 37 56 64 18 178 188 24 187 39 13 286 1.6 31 9.1

CAL YR 1998 TOTAL 57187.13 MEAN 158 WAX 5368 WIN 15 AC-FT 113388 CFSW 3.98 IN 52.98 WTR YR 1991 TOTAL 41649.93 MEAN 114 WAX 3248 WIN 68 AC-FT 82618 CFSW 2.85 IN 38.64

