

Effects of a natural precipitation gradient on fish and macroinvertebrate assemblages

Sean Kinard Corresp., 1, Christopher Patrick 1, Fernando Carvallo 2

Corresponding Author: Sean Kinard Email address: skkinard@vims.edu

In the American Southwest, conditions are expected to become more arid. To enhance our understanding of how freshwater communities will adjust to these shifts in water-cycle dynamics, we employed a space for time approach along a precipitation gradient from semi-arid to sub-humid on the Texas Coastal Prairie. In the Spring of 2017, we conducted surveys of 10 USGS gauged, wadeable streams spanning a natural precipitation gradient; we measured nutrients, water chemistry, habitat characteristics, benthic macroinvertebrates, and fish community data. We also observed a positive relationship between fish diversity and mean annual rainfall, conductivity and surface runoff. Macroinvertebrate diversity did not correlate with annual precipitation but was correlated with low flow pulse percent. The compositional shifts of fish and invertebrate communities along the gradient indicate both top-down and bottom-up controls on community assembly. Semi-arid sites contain euryhaline, and rapid proliferating taxa. Sub-humid sites contain migratory euryhaline fish and fish predators which impose top-down controls on primary consumers. Proceeding from humid to arid, low-flow conditions (high solute concentrations and habitat fragmentation) restrict fish compositions. These results indicate that small future changes in precipitation regime in this region may result in abrupt transitions into new community states.

¹ Department of Biological Sciences, Virginia Institute of Marine Science, Gloucester Point, Virginia, United States of America

² Department of Life Sciences, Texas A&M Corpus Christi, Corpus Christi, Texas, United States of America



40 41

1 2 3	Manuscript Title Effects of a natural precipitation gradient on fish and macroinvertebrate assemblages
4 5 6	Sean Kinard ¹ , Christopher Patrick ¹ , Fernando Carvallo ²
7 8 9 10	 Virginia Institute of Marine Science 1370 Greate Road, Gloucester Point, Virginia 23062-1346 Texas A&M Corpus Christi 6300 Ocean Dr, Corpus Christi, TX 78412
12 13 14 15 16	Corresponding Author: Sean Kinard ¹ 6528 Quail Hollow Dr, Hayes VA 23072, USA Email address: skkinard@vims.edu
17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33	Abstract: In the American Southwest, conditions are expected to become more arid. To enhance our understanding of how freshwater communities will adjust to these shifts in water-cycle dynamics, we employed a space for time approach along a precipitation gradient from semi-arid to sub-humid on the Texas Coastal Prairie. In the Spring of 2017, we conducted surveys of 10 USGS gauged, wadeable streams spanning a natural precipitation gradient; we measured nutrients, water chemistry, habitat characteristics, benthic macroinvertebrates, and fish community data. We also observed a positive relationship between fish diversity and mean annual rainfall (<i>p</i> -value = 0.008), conductivity (<i>p</i> -value = 0.048) and surface runoff (<i>p</i> -value = 0.002). Macroinvertebrate diversity did not correlate with annual precipitation but was correlated with low flow pulse percent (<i>p</i> -value = 0.046). The compositional shifts of fish and invertebrate communities along the gradient indicate both top-down and bottom-up controls on community assembly. Semi-arid sites contain euryhaline, and rapid proliferating taxa. Sub-humid sites contain migratory euryhaline fish and fish predators which impose top-down controls on primary consumers. Proceeding from humid to arid, low-flow conditions (high solute concentrations and habitat fragmentation) restrict fish compositions. These results indicate that small future changes in precipitation regime in this region may result in abrupt transitions into new community states.
34 35 36 37 38	Introduction: Anthropogenic climate change creates an urgent need to understand the relationship between biological communities and climate (Wrona, Prowse et al. 2006). A warmer, more energetic atmosphere intensifies the hydrological cycle (i.e. patterns of precipitation and evaporation), causing wet regions to become wetter and dry regions become drier (Allen and

42 systems contain many species with limited dispersal capabilities which are highly sensitive to 43

changes water temperature and availability (Woodward, Perkins et al. 2010). Global hydrological 44

Ingram 2002), and increases the frequency and intensity of extreme weather events (Held and

ecosystems, especially in arid and semi-arid regions (Grimm, Chapin et al. 2013). Freshwater

Soden 2006). The predicted changes in precipitation patterns will have significant effects on

models predict region-specific changes in annual flow regime including shifts from intermittent



to perennial streamflow in Arizona, New Mexico, and West Texas. However, it is unclear how the biological communities within the stream ecosystems will respond to the predicted changes to the hydrologic cycle. Therefore, clarifying mechanistic links between climate drivers and instream biological communities will improve our ability to predict to the effects of anthropogenic climate change on lotic ecosystems.

Streams ecosystems are shaped by flow regimes which regulate the physical extent of aquatic habitat, the water quality, sourcing and exchange rates of material, habitat connectivity and diversity (Rolls, Leigh et al. 2012). In addition to streamflow mechanisms, streamside vegetation mediates interactions with watershed nutrients, carbon and light inputs to streams (Schade, G. Fisher et al. 2001). Precipitation regime is the primary regulator of both streamflow and riparian characteristics. With predicted changes in flood and drought characteristics under global warming (Hirabayashi, Kanae et al. 2008), it is imperative to understand the mechanistic links between precipitation, streamflow, and riparian interactions with aquatic biological communities

Hierarchical community assembly models can help us organize our hypotheses regarding impacts of climate change on stream communities (Poff 1997). Assuming organisms can disperse to a habitat, they must be able to survive in the local environment (abiotic filters) and successfully reproduce in the presence of other organisms exerting pressures (biotic interactions) such as competition and predation (Patrick and Swan 2011). Abiotic filters are conceptually easy to understand. Species have physiological tolerances which limit their distribution across environmental gradients (Whittaker, Willis et al. 2001), if climate change alters those gradients we can expect concordant changes in species distributions. However, understanding how the environment affects biotic interactions is more challenging due to the complex sets of interactions that govern these processes (Seabra, Wethey et al. 2015). As a result, our understanding of the role of environmental filters on community assembly is disjointed due to the vastly different spatial scales of typical biogeographical and community ecology studies (Ricklefs and Jenkins 2011).

Observational surveys of existing communities spatially distributed along environmental gradients can be used in a space-for-time substitution to infer how communities will change through time as environmental conditions shift (Ricklefs and Jenkins 2011). The approach allows for links to be drawn between climate drivers, local environmental conditions, and organism abundances. Species co-occurrence patterns along environmental gradients can also shed light on possible shifts in biotic interactions (D'Amen, Mod et al. 2018). However, the space-for-time substitution approach assumes that observed ecological differences along the spatial gradient are the sole product of corresponding changes in climate. This assumption may be unfair given that biogeographical studies have revealed that dispersal limitation, habitat heterogeneity, and local evolution can also contribute to current spatial patterns in community composition (Jacob et al. 2015). These studies are typically large in scale, covering vast distances (thousands of km) in order to capture climate gradients. These large scales make the precise mechanisms for observed biological changes difficult to ascertain due to covarying environmental variables (e.g., elevation, geology, human impacts). Thus, while current literature demonstrates that biome shifts occur across temperature and latitudinal gradients (De Frenne, Graae et al. 2013), the value of these observational studies for forecasting community responses to climate change is hindered by the many confounding variables. The power of using the space-for-time approach to delineate the intricacies of hydrologic cycle-ecosystem relationships is enhanced in study systems with limited



confounding environmental variables (i.e. temperature, elevation, distance, and underlying geology).

Fortunately, the Texas Coastal Prairie (TCP) within the Western Gulf coastal grasslands is an ideal system evaluating the effect of hydrologic climate change on ecological communities. The Western Gulf coastal grasslands are a subtropical ecotone that spans Louisiana, Texas, and northern Mexico's coastal areas. From east to west to climate becomes more arid, with gradual change for much of the coast and a region of rapid change located in southern Texas. In this region the annual rainfall changes from 55cm•yr-1 (semi-arid) to 135 cm•yr-1 (sub-humid) over a 300 km gradient (Falcone 2011), but there are minimal changes in elevation, air temperature, underlying geology, and human land use. Thus, studying natural ecosystems that span the TCP maximizes our ability to detect relationships between annual precipitation and ecosystem processes in the absence of covarying factors.

We used a section of the TCP where precipitation changes most quickly as a model system to evaluate how changes in precipitation alter stream communities. As conditions become wetter, there is an observable ecological shift from Thornwood groves in the semi-arid West to Live oak forests towards the East (Chapman BR 2018). In addition to its value as a case study region, there is limited prior biological sampling by state and federal agencies of running waters in the TCP, so sampling efforts enhance our understanding of subtropical ecosystems (US EPA 2016). Along the rainfall gradient we surveyed 10 USGS gauged wadeable streams for fish, benthic macroinvertebrates, and environmental variables. Our objectives were to: 1) Identify patterns in the diversity and composition of fish and macroinvertebrates communities that correspond to changes in precipitation, and 2) identify environmental drivers that mediate the effects of climate on community processes. We expected that annual precipitation would be positively correlated with community diversity because humid precipitation regimes are expected to create more stable environmental conditions by creating habitat heterogeneity and predictable flow regimes which promote the development of greater biodiversity (Boulton, Peterson et al. 1992). We further expected that evapotranspiration by riparian vegetation would increase solute concentrations in semi-arid streams, particularly during base flows (Tabacchi, Lambs et al. 2000, Lupon, Bernal et al. 2016), creating environmental filters that limit recruitment of sensitive fish and macroinvertebrates (hereafter referred to as invertebrates).

Methods

Study Region: The Texas Coastal Prairie contains grassland prairie with forested areas occurring primarily along riverine systems. During March and April of 2017, we sampled ten, wadable, perennial streams which span 12 counties from Kleberg County to Montgomery in South-Central Texas, USA (Fig. 1). Each study site is located within 100 meters of a USGS stream gauge which continuously monitor streamflow and climate data year-round. Study sites were chosen to maximize differences in precipitation with minimal changes in underlying geology and elevation. The annual precipitation ranges from 61-121 cm within the study region which spans a linear distance from end to end of 378 km (Falcone 2011). The surface geology is characterized by fine clays, quaternary and sedimentary sand. The streams have similar elevations (14-62 m), substrates (quaternary), and average air temperatures (19.8-22.1°C) (Appendix-site). Sampling was conducted by students and faculty at Texas A&M (Corpus Christi) under permit SPR-0716-170, granted by Texas Parks and Wildlife Department.

Biological Sampling: Fish communities were sampled using a Smith-Root LR-24 Backpack in a single pass survey of a 100-meter reach (Lamberti 2007). The reach length was approximately 25 times the average stream width (4.1m), in accordance with EPA rapid bioassessment protocols (Barbour, Gerritsen et al. 1999). Study sites are characterized by low variation in geomorphology and overall habitat heterogeneity resulting in high success in assessing community composition over a shorter distance. Fish species were field identified to species using a field guide (Thomas C 2007) and photographed. Several specimens of each species were euthanized using tricaine mesylate (MS-222) and stored in >70% denatured ethanol as voucher specimens for lab confirmation of species identification. Fish Voucher specimens were identified using the Texas Academy of Science dichotomous key (Hubbs 2008) and cross referenced with field identifications. Vertebrate sampling was permitted by the Institutional Animal Care and Use Committee, Texas A&M University Corpus Christi (AUP# 05-17).

Invertebrates were collected using a 0.305m wide D-frame net equipped with 500-µm mesh. Twenty 0.093 m² samples were collected via a combination of kick and sweep sampling from a representative distribution of best available habitat (riffles, large woody debris, overhanging vegetation). Samples were pooled and field rinsed in a 500-µm sieve bucket. After removal of rinsed larger sticks and leaves, the entire sample was preserved with the addition of 95% EtOH for transport to the lab. In the lab, samples were spread across a gridded sampling tray and randomly selected grid cells were picked to completion until the total count was > 300 individuals. Samples containing less than 300 individuals were picked to completion. Invertebrates were identified to lowest taxonomic resolution (typically genus) using taxonomic keys cross referenced with species observations recorded by the TCEQ's (Texas Commission on Environmental Quality) Surface Water Quality Monitoring Program (Wiggins 2015, Merritt, Cummins et al. 2019). The sum of individuals in each taxon were multiplied by the fraction of unpicked sample and reported as abundance of individuals per square meter.

Environmental Data: For each stream, we averaged values for each of the following habitat measurements that were taken at 4 cross-sections spaced 25m apart. Canopy cover was measured using a spherical densiometer. A Rosgen Index value was calculated by dividing the bank-full width by the maximum depth (DL 2001). Bank height was recorded as vertical difference between water level and the height of the first bench. We estimated Sediment grain size within each cross-section using Wentworth size categories to calculate a median grain-size (d50) (Wentworth 1922). Oxygen, temperature (T_{water}), conductivity, turbidity, and pH were measured at each point using a YSI ProDSS multiparameter probe. Two 60 mL water samples were collected and filtered through a pre-combusted (500°C for 4 hours) glass fiber filter (Whatman GF/F) into acid washed amber bottles, transferred to the lab in a cooler on ice, and stored frozen (-20°C) until analysis for nutrients (NH₄⁺, NO₃⁻, and PO₄⁻). Water samples were run using colorimetric methods on a latchet autoanalyzer by the Oklahoma University Soil Water and Forage Laboratory.

In addition to the habitat metrics measured in the field, we mined climate and watershed data (average annual precipitation (AP), relative humidity (RH), mean-annual potential evapotranspiration (PET), proportion of forested riparian zone (Rip.forest), and proportions of watershed forest (Bas.forest), agriculture (Bas.plant), and urban development (Bas.dev), soil permeability (Soil.Perm), soil organic content (Soil.Org), and runoff factor) from the US Geologic Surveyors Geospatial Attributes of Gages for Evaluating Streamflow, version II dataset (Falcone 2011). A twenty-year continuous daily flow record was downloaded for each site

 \bigcirc



(except Tranquitas Creek which only had 4 years of available data) from the USGS water services (Falcone 2011).

In order to evaluate the typical flow regime of each site in the context of seasonal droughts, floods, and overall variation in flow, we calculated the following four variables: average daily discharge (Discharge), the flashiness index (Flash Index = cumulative changes in day to day daily flow / cumulative flow), the high flow Pulse Percentage (HFPP = Percent of time daily flow is above 3 times the median daily flow), and the low flow pulse percentage (LFPP = times where daily discharge drops below the 25th percentile) (Olden and Poff 2003, Patrick and Yuan 2017).

Analyses: For each community (fish, invertebrates) we calculated Shannon diversity and rarified taxonomic richness (Hurlbert 1971). Diversity and richness measures were calculated using the Vegan Library (Oksanen, Blanchet et al. 2019) in the statistical program R (R Core Team 2019). To evaluate the effect of rainfall on environmental variables that may influence the biota, we used linear regression to examine relationships between environmental variables and annual precipitation. We then used linear regression to evaluate the effect rainfall and environmental variables on fish and invertebrate community metrics. Prior to analyses conductivity and NO₃ concentrations were natural log transformed to satisfy the test assumption of normality.



To evaluate how community composition changed along the gradient, we ordinated each taxa group (fish, invertebrates) across sites using non-metric multidimensional scaling (NMDS) using the 'metaMDS' within the VEGAN R package. The function runs NMDS with multiple starting configurations (n=100), compares results, and stops after finding a similar minimum stress solution. The minimum stress solution is scaled, rotated, and then species scores are added to the configuration as weighted averages (Oksanen, Blanchet et al. 2019). Using the climate data, sites were grouped using Ward's minimum variance method for hierarchical clustering (Ward 1963). Finally, we fit environmental variables to each ordination using the 'envfit' function within the VEGAN R package; this function fits environmental vectors onto the ordination and calculates the maximum correlation with the projection of points (sites in this case). To ease in visual interpretation of the plots we only retained environmental variables that were significant at an $\alpha > 0.10$ and explained > 40% of the variation. Retained environmental vectors were overlaid on top of the NMDS plots. The direction of the vector indicates the axis of the variable within the ordination space and the length of the arrow is proportional to its correlation with the projected points (Oksanen, Blanchet et al. 2019).

Results

Site Overview: Regression analysis of 26 environmental variables indicates 20 significant (R² > 0.40 and a p-value < 0.05) relationships reported in Appendix-regressions. We observed several significant relationships between mean annual precipitation and measured environmental variables (Appendix-regression). Surface runoff was positive related with, whereas conductivity and potential evaporation were negatively related to precipitation (Fig. 2). Increasing PET is correlated with increases in T_{water} and decreases in Soil.Org. Lastly, relative humidity correlates negatively with LFPP.

- 223 Significant environmental regressions excluding climate variables are summarized as follows
- 224 (Appendix-regression); HFPP correlates negatively with PO₄-, canopy coverage is positively
- 225 correlated with NH₄⁺, Rosgen Index is positively correlated with Soil.Org and turbidity, bank
- 226 height is positively correlated with Soil.Perm, pH correlates positively with conductivity,



turbidity is positively correlated with NO₃⁻ and Soil.Org, conductivity is positively colinear with pH and is negatively correlated with AP and runoff factor, NO₃⁻ correlates positively with turbidity and Soil.Org. Bas.dev positively correlates with Soil.Org. Bas.plant correlates negatively with Soil.Perm.

Fish Community: In total, 18 fish species were identified within the surveyed sites. Proceeding from semi-arid to sub-humid sites, fish Shannon index increases from 0.64 - 1.81, richness increases from 2 - 7 species, and rarified richness increases from 2.09 - 5.48 species. Rarified Richness_{fish} increases with increasing Rip.forest (R²=0.404, p=0.048) or runoff factor (R²=0.415, p=0.044). Shannon Index_{fish} decreases with increasing PET (R²=0.518, p=0.019), conductivity (R²=0.406, p=0.048), or NH₄⁺ (R²=0.445, p=0.035). Shannon Index_{fish} increases with increasing AP (R²=0.602, p=0.019) and runoff factor (R²=0.716, p=0.002).

Fish abundances are found in Appendix-fish and are summarized as follows. Fish species found throughout the study region include, Red Shiner (*Cyprinella lutrensis*), Western Mosquitofish (*Gambusia affinis*), Longear Sunfish (*Lepomis megalotis*), and Bullhead minnow (*Pimephales vigilax*). Sailfin molly (*Poecilia latipinna*) was found only in Western (semi-arid) sites. Several fish species were found only in mesic sites including Rio Grande cichlid (*Herichthys cyanoguttatus*) and slough darter (*Etheostoma gracile*). Fish species found throughout the central and Eastern sites (mesic and sub-humid) include blackstripe topminnow (Fundulus notatus), green sunfish (*Lepomis cyanellus*), warmouth sunfish (*Lepomis gulosus*), bluegill sunfish (*Lepomis macrochirus*), and dollar sunfish (*Lepomis marginatus*). Fish species found only in the Eastern (sub-humid) sites include black bullhead catfish (*Ameiurus melas*), American eel (*Anguilla rostrata*), blacktail shiner (*Cyprinella venusta*), redbreast sunfish (*Lepomis auritus*), and orangespotted sunfish (*Lepomis humilis*).

The best solution for the NMDS ordination of fish community data had a stress value of 0.156 indicating a good fit of the data (Oksanen et al. 2019). The NMDS ordination of fish assemblages indicate compositional shifts across the precipitation gradient (Fig. 3). Hierarchical clustering resulted in 3 site groupings labeled "semi-arid" (yellow), "mesic" (green), and "sub-humid" (blue). Semi-arid fish communities are dominated by sailfin molly, western mosquitofish, and bullhead minnow. Fish communities in mesic and sub-humid climates contain a variety of sunfish species, but mesic streams uniquely contain Rio Grande cichlid. Sub-humid streams uniquely contain hogchoker (*Trinectes maculatus*), black bullhead catfish, and blacktail shiner. Significant fitted environmental variables on fish community NMDS include RH and LFPP (Appendix-ordination).

Invertebrate Community: A total of 94 invertebrate genera were identified within the study region. Invertebrate richness ranges 7–29 genera with the highest values (29, 26, and 27) occurring at three mesic sites (Aransas, Perdido and Mission respectively). Invertebrate Shannon index ranges 1.83–3.30 with higher values (3.30, 3.18 and 3.28) at three mesic sites (Aransas, Perdido, and Mission respectively) (Appendix-invert). Rarified richness_{invertebrate} did not correlate significantly with environmental predictors. Shannon Index_{invertebrate} decreases with increasing LFPP (R²=0.411, p-value=0.046). Since maximum richness and Shannon index values were observed at sites in the middle of the precipitation gradient, we conducted a second order quadratic regression between Shannon Index_{invertebrate} and AP (R²=0.319, p-value=0.260) (Fig. 2).

Invertebrate community abundances are found in Appendix-invert and are summarized as follows. We identified 17 genera of coleoptera; mesic and sub-humid sites contained a variety,



but the semi-arid sites (Tranquitas and San Fernando) contain 0-1 genera (Stenelmis). We identified 17 genera of Ephemeroptera throughout the study region; no Ephemeroptera were identified at two semi-arid sites (Tranquitas and San Fernando) and one sub-humid site (Placedo). We identified 17 genera of Gastropoda within the study region; abundances are relatively high (exceeding 1000 per m²) for several genera in the semi-arid and mesic sites (Tranquitas, San Fernando, and Aransas). 16 genera of Hemiptera were identified across the region; none were found in one semi-arid (Tranquitas) and three sub-humid sites (Placedo, Garcitas, and Bear Branch). 10 genera of Odonata were identified only in one semi-arid (San Fernando) and four mesic sites (Aransas, Medio, Perdido, and Mission). 10 genera of Trichoptera were identified across the study region, but one semi-arid site (Tranquitas) had none. 2 genera of Amphipoda were identified with Hyalella occurring at nine sites (absent at Tranquitas) and Gammarus only occurring at two mesic sites (San Fernando and Aransas). 2 genera of Bivalvia were identified with Corbicula occurring at six sites across the region and Pisidium only occurring at Placedo. 2 genera of Decapoda were identified with Palaemonetes occurring at nine sites (absent at Tranquitas) and Orconectes occurring at Aransas, 1 genus of Isopoda was identified (Caecidotea) and was only present at San Fernando.

The best solution for the NMDS ordination of the invertebrate community data had a stress value of 0.098 indicating a good fit of the data (Oksanen 2013). The NMDS ordination of invertebrate assemblages (Fig. 3) displays compositional shifts along the precipitation gradient. Invertebrate assemblages in semi-arid climate contain a variety of gastropod taxa including *Amnicola sp. Bythinia sp. And melanoides sp.* Mesic communities contain species from a greater number of taxonomic orders including Ephemeroptera, Trichoptera, Coleoptera, and Hemiptera. Sub-humid communities contain a greater proportion of Crustaceans including *Palaemonetes sp.*, *Orconectes sp.*, and isopods in the genus *Caecidotea*. Significant fitted environmental variables on invertebrate community NMDS include RH and LFPP, and conductivity (Appendixordination).

Discussion

Using the Texas Coastal Prairie (TCP) as a model system, our goal was to quantify patterns in the diversity and composition of stream communities along an extreme precipitation gradient to better understanding how streams might respond to future changes in mean annual rainfall. Our observational study identified strong compositional shifts in both fish and invertebrate communities along the precipitation gradient. We also observed a positive relationship between fish diversity and mean annual rainfall, matching expectations, however, invertebrate diversity did not exhibit the expected relationships with rainfall. Environmental data collected at each site suggest several mechanistic drivers of these changes operating through water solute concentrations and flow regimes. Below we discuss these results, place in the context of other literature, and make suggestions for future work.

The lack of observed relationships between annual precipitation (AP) and most environmental variables supports the assertion that TCP is an exemplary region to conduct space for time substitutions to make useful ecological predictions regarding climate change. While we did observe relationships between AP, potential evapotranspiration (PET), and runoff and water quality variables such as conductivity and nutrients as well as riparian cover, these relationships are likely causal and important mechanistic pieces of the relationships between AP and stream communities. The field-measured riparian data (canopy) proved uninformative due to outlier effects brought on by sub-urban floodway maintenance at our most humid site, Bear Creek. So,



we restrict our discussion of riparian-effects on community assembly to the watershed-level metric (Bas.forest), supplied by the USGS.

The fish communities displayed a pattern of increasing diversity, rarified richness, and compositional turnover moving from the drier to wetter sides of the survey region (Fig. 2). The wetter sites were characterized by an increase in the diversity of sunfishes and the addition of several marine migrants including hogchoker and American eel. These compositional shifts connect with quantitative relationships between environmental variables and diversity, suggesting mechanistic pathways through which precipitation is structuring the stream communities. As precipitation increases, fish communities structure diversifies to include competitive omnivores and predators. Mesic sites contain a plurality of centrarchids; species with 3-7 year lifespans, annual breeding, nesting strategies, and are omnivores (Cooke and Philipp 2009).

Sub-humid sites contain larger predator taxa including catfish, largemouth bass, warmouth sunfish, and green sunfish. Most of these species are ambush predators that reside within alcoves and woody debris, consuming a mixture of insects and small fish. These same taxa likely benefited from rainfall via an indirect effect on riparian vegetation. The relationship between canopy cover and rainfall was positive but non-significant, but Rip. forest had a strong positive relationship with fish diversity (Fig. 2). Mechanistically, riparian trees provide appropriate conditions for fish taxa via root-stabilized undercut banks or large woody debris within the channel (Krzeminska, Kerkhof et al. 2019). Large wood and bank stabilization are particularly important in these grassland prairie streams because the substrate is largely unconsolidated sand and there is little natural structure. Although not accounted for in this study, the changes in riparian vegetation from grasses to trees across this precipitation gradient may correspond to shift from autochthonous production to allochthonous production, fundamentally changing the basal resources within these aquatic systems (Hagen, McTammany et al. 2010). Further investigation of these mechanisms will require woody debris counts, bank characterizations, and stable isotope analysis of food web resources.

On the other extreme, communities in semi-arid streams (<75 cm annual precipitation) were composed of rugged species able to tolerate high salinities including Sailfin Molly (95 psu) and Western Mosquitofish (58.5 psu) (Page and Burr, B.M. 1991). The observed negative relationship between AP and conductivity, and conductivity and fish diversity point to the importance of rainfall creating low-flow hydrologic conditions (Fig. 2). Aquatic ecosystems in arid regions are prone to salinization (Williams 2002), which in conjunction with evapotranspiration and presence of agriculture contribute to base flow salinity concentrations that limit diversity by permitting only species with specialized osmoregulatory mechanisms (East, Wilcut et al. 2017). Taken together, our results indicate that regional decreasing AP restricts fish community assembly by increasing osmoregulatory stress in aquatic vertebrates.

In addition to poor water quality conditions, low-flow hydrologic regimes can limit habitat connectivity and diversity. Migratory, euryhaline species including Rio Grande cichlid (0-27.5 PSU), hogchoker (1-30 PSU), and American eel (1-36 PSU) are only found in Mesic and Sub-Humid streams (Fig. 3). Hogchoker typically reside in brackish estuaries (1-25psu) and make seasonal migrations upstream to spawn (Koski 1978). American Eel are catadromous species in which adults migrate to the Sargasso Sea to reproduce and juvenile migrate upstream to rear (Wenner 1978). Rio Grande Cichlid seek thermal refugia in deeper pools or estuaries during the winter months until temperatures rise and flows permit dispersal in late Spring (Rehage, Blanchard et al. 2016). Given the similar distances to nearby estuaries, we suspect

anadromous, euryhaline taxa are excluded from semi-arid streams due to increased habitat fragmentation and the unpredictability of freshets in semi-arid climate. In order to substantiate the claim that low-flow hydrology restricts fish movement in the semi-arid region of the precipitation gradient, we need to conduct seasonal surveys during wet and dry seasons.

While the invertebrate communities showed compositional shifts along the precipitation gradient, unlike fish, there was not a positive relationship with Shannon diversity. Many invertebrate taxa can mitigate the effects of drought-induced habitat fragmentation by seeking refuge in the hyporheic zone, interstitial spaces, and desiccation-resistant life-stages (Boulton, Peterson et al. 1992, Boulton 2003). The lack of diversity trends across the precipitation gradient could be attributed to the inherently larger regional pool or invertebrate species, many of which can tolerate drought conditions that typify the semi-arid region of the precipitation gradient.

Low flow pulse percentage (LFPP) was the sole significant predictor of invertebrate community diversity (Fig. 2). LFPP increases as RH decreases and indicates the prevalence of low-flow hydrologic regime in the semi-arid region of the precipitation gradient. Additionally, invertebrate community compositions shift along the precipitation gradient and these shifts coincide with changes in LFPP (Fig. 3). Semi-arid streams communities are composed of grazing Gastropoda and some predatory Hemiptera. Red-rimmed Melania (*Melanoides tuberculata*), dominate these systems, likely due to their salinity tolerance (0-23 PSU) and their ability to rapidly proliferate following dewatering due to their short life cycles, reaching sexual maturity within 21-62 days (Krumholz 1948, Farani, Nogueira et al. 2015).

On the other extreme of the gradient, sub-humid sites contain an abundance of grazer and omnivore taxa including Ephemeroptera, Amphipoda, and Trichopteran. Between these regions, mesic sites contain a mixture of grazers, omnivores and an abundance of predators including Odonata and Hemiptera. As precipitation increases, there is a shift in primary consumers from Gastropoda to Ephemerpotera and Trichoptera; the latter species indicating enhance water quality conditions (Rosenberg and Resh 1993). Additionally, these taxa have longer life cycles (26-261days) than the prominent grazers at semi-arid sites (Jackson and Sweeney 1995). These results indicate that drought also acts to regulate insect communities at the primary consumer level. Discerning the specific mechanisms of drought (intensity, duration, seasonality, and predictability) will require continuous sampling across the precipitation gradient.

Initially, the maximum invertebrate diversity in mesic sites can be attributed to overlapping dispersal from the extreme climate regions. However, the compositional shifts broadly indicate that predation and competition play larger roles in community assembly at mesic and sub-humid sites. Specifically, the distribution of fish predators has large top-down controls on invertebrate community dynamics (Dahl and Greenberg 1998). Here, we believe fish are superior insectivores compared to Hemiptera and Odonata and that fish predation at sub-humid sites restricts invertebrate communities to species with anti-predator adaptations including small size, sedentary forage strategies, and armoring (Straile and Hälbich 2000). Our results suggest that as conditions become more arid, top-down regulation by fish predators is reduced resulting in a proliferation of insect predators in mesic and semi-arid streams. A more thorough invertebrate community analysis of primary consumers will include sediment core sampling and functional trait analysis.

Acknowledgements



409 Jennifer Whitt and Ian Whitt 410 411 **References:** 412 Allen, M. R. and W. J. Ingram (2002). "Constraints on future changes in climate and the 413 hydrologic cycle." Nature 419(6903): 224-+. 414 415 Barbour, M. T., J. Gerritsen and J. B. S. B.D. Snyder (1999). Rapid Bioassessment Protocols for 416 Use in Streams and Wadeable Rivers: Periphyton, Benthic Macroinvertebrates and Fish, Second 417 Edition. Washington, D.C., U.S. Environmental Protection Agency; Office of Water. 418 419 Boulton, A. J. (2003). "Parallels and contrasts in the effects of drought on stream 420 macroinvertebrate assemblages." Freshwater Biology **48**(7): 1173-1185. 421 422 Boulton, A. J., C. G. Peterson, N. B. Grimm and S. G. Fisher (1992). "Stability of an Aquatic 423 Macroinvertebrate Community in a Multiyear Hydrologic Disturbance Regime." Ecology **73**(6): 424 2192-2207. 425 426 Chapman BR, B. E. (2018). The Natural History of Texas. College Station, Texas A&M 427 University Press. 428 429 Cooke, S. and D. Philipp (2009). Centrarchid Fishes: Diversity, Biology, and Conservation. 430 D'Amen, M., H. K. Mod, N. J. Gotelli and A. Guisan (2018). "Disentangling biotic interactions, 431 environmental filters, and dispersal limitation as drivers of species co-occurrence." Ecography 432 **41**(8): 1233-1244. 433 434 Dahl, J. and L. A. Greenberg (1998). "Effects of fish predation and habitat type on stream 435 benthic communities." Hydrobiologia **361**: 67-76. 436 437 De Frenne, P., B. J. Graae, F. Rodriguez-Sanchez, A. Kolb, O. Chabrerie, G. Decocq, H. De 438 Kort, A. De Schrijver, M. Diekmann, O. Eriksson, R. Gruwez, M. Hermy, J. Lenoir, J. Plue, D. 439 A. Coomes and K. Verheyen (2013). "Latitudinal gradients as natural laboratories to infer 440 species' responses to temperature." Journal of Ecology 101(3): 784-795. 441 442 DL, R. (2001). "A Stream Channel Assessment Methodology; Proceedings of 7th Federal 443 Interagency Sedimentation Conference." 26. 444 East, J. L., C. Wilcut and A. A. Pease (2017). "Aquatic food-web structure along a salinized 445 446 dryland river." Freshwater Biology 62(4): 681-694. 447 448 Falcone, J. (2011). GAGES-II: Geospatial Attributes of Gauges for Evaluating Streamflow. 449 Reston, Virginia, U.S. Geological Survey.



- 451 Farani, G. L., M. Nogueira, R. Johnsson and E. Neves (2015). "The salt tolerance of the
- 452 freshwater snail Melanoides tuberculata (Mollusca, Gastropoda), a bioinvader gastropod." 10:
- 453 212-221.
- 454 Grimm, N. B., F. S. Chapin, B. Bierwagen, P. Gonzalez and P. M. Groffman (2013). "The
- 455 impacts of climate change on ecosystem structure and function." Frontiers in ecology and the
- 456 environment **11**(9): 474-482.

- Hagen, E. M., M. E. McTammany, J. R. Webster and E. F. Benfield (2010). "Shifts in
- allochthonous input and autochthonous production in streams along an agricultural land-use
- 460 gradient." Hydrobiologia **655**(1): 61-77.

461

Held, I. M. and B. J. Soden (2006). "Robust responses of the hydrological cycle to global warming." Journal of Climate **19**(21): 5686-5699.

464

- Hirabayashi, Y., S. Kanae, S. Emori, T. Oki and M. Kimoto (2008). "Global projections of
- changing risks of floods and droughts in a changing climate." <u>Hydrological Sciences Journal</u>
- **467 53**(4): 754-772.

468

- Hubbs, C. (2008). An Annotated Checklist of the Freshwater Fishes of Texas, with Keys to
- 470 Identification of Species. R. J. E. a. G. P. Garrett, Texas Academy of Science.

471

- Jackson, J. K. and B. W. Sweeney (1995). "Egg and Larval Development Times For 35 Species
- 473 of Tropical Stream Insects from Costa-Rica." <u>Journal of the North American Benthological</u>
- 474 Society **14**(1): 115-130.

475

- 476 Koski, R. T. (1978). "Age, Growth, and Maturity of Hogchoker, *Trinectes* maculatus, In Hudson
- 477 River, New York." Transactions of the American Fisheries Society **107**(3): 449-453.

478

- Krumholz, L. A. (1948). "Reproduction in the Western Mosquitofish, Gambusia-Affinis-Affinis
- 480 (Baird and Girard), and Its Use in Mosquito Control." Ecological Monographs 18(1): 1-43.

481

- 482 Krzeminska, D., T. Kerkhof, K. Skaalsveen and J. Stolte (2019). "Effect of riparian vegetation
- on stream bank stability in small agricultural catchments." Catena 172: 87-96.

484

- Lamberti, H. (2007). Methods in Stream Ecology, Elsevier Inc.
- 486 Lupon, A., S. Bernal, S. Poblador, E. Marti and F. Sabater (2016). "The influence of riparian
- 487 evapotranspiration on stream hydrology and nitrogen retention in a subhumid Mediterranean
- catchment." Hydrology and Earth System Sciences **20**(9): 3831-3842.

489

- 490 Merritt, R. W., K. W. Cummins and M. B. Berg (2019). An introduction to the Aquatic insects of
- 491 North America. Dubuque, Iowa, Kendall/Hunt Pub. Co.

492

- Oksanen, J., F. G. Blanchet, M. Friendly, R. Kindt, P. Legendre, D. McGlinn, P. R. Minchin, R.
- 494 B. O'Hara, G. L. Simpson, P. Solymos, M. Henry, H. Stevens, E. Szoecs and H. Wagner (2019).
- 495 "vegan: Community Ecology Package."



- Olden, J. D. and N. L. Poff (2003). "Toward a mechanistic understanding and prediction of biotic
- 498 homogenization." American Naturalist **162**(4): 442-460.

- Patrick, C. J. and C. M. Swan (2011). "Reconstructing the assembly of a stream-insect
- metacommunity." Journal of the North American Benthological Society **30**(1): 259-272.

502

- Patrick, C. J. and L. L. Yuan (2017). "Modeled hydrologic metrics show links between
- 504 hydrology and the functional composition of stream assemblages." Ecological Applications
- 505 **27**(5): 1605-1617.

506

- 507 Poff, N. L. (1997). "Landscape filters and species traits: Towards mechanistic understanding and
- prediction in stream ecology." Journal of the North American Benthological Society 16(2): 391-
- 509 409.

510

- R Core Team (2019). R: A Language and Environment for Statistical Computing. Vienna,
- Austria, R Foundation for Statistical Computing.

513

- Rehage, J. S., J. R. Blanchard, R. E. Boucek, J. J. Lorenz and M. Robinson (2016). "Knocking
- back invasions: variable resistance and resilience to multiple cold spells in native vs. nonnative
- 516 fishes." Ecosphere **7**(6): 13.

517

- Ricklefs, R. E. and D. G. Jenkins (2011). "Biogeography and ecology: towards the integration of
- 519 two disciplines." Philosophical Transactions of the Royal Society B-Biological Sciences
- 520 **366**(1576): 2438-2448.

521

- Rolls, R. J., C. Leigh and F. Sheldon (2012). "Mechanistic effects of low-flow hydrology on
- 523 riverine ecosystems: ecological principles and consequences of alteration." Freshwater Science
- 524 **31**(4): 1163-1186.

525

- S26 Rosenberg, D. M. and V. H. Resh (1993). Freshwater Biomonitoring and Benthic
- 527 Macroinvertebrates. Boston, Kluwer Academic Publishers.

528

- 529 Schade, J., S. G. Fisher, N. Grimm and J. A. Seddon (2001). The Influence of Riparian Shrub on
- 530 Nitrogen Cycling in a Sonoran Desert Stream.

531

- Seabra, R., D. S. Wethey, A. M. Santos and F. P. Lima (2015). "Understanding complex
- 533 biogeographic responses to climate change." Scientific Reports 5.

534

- 535 Straile, D. and A. Hälbich (2000). "Life History and Multiple Antipredator Defenses of an
- 536 Invertebrate Pelagic Predator, Bythotrephes longimanus." Ecology 81: 150-163.

537

- Tabacchi, E., L. Lambs, H. Guilloy, A. M. Planty-Tabacchi, E. Muller and H. Decamps (2000).
- "Impacts of riparian vegetation on hydrological processes." Hydrological Processes **14**(16-17):
- 540 2959-2976.





- 542 Thomas C, B. T., Whiteside BG (2007). A Field Guide: Freshwater Fishes of Texas. College
- 543 Station, Texas, Texas A&M University Press.

- Ward, J. H. (1963). "Hierarchical Grouping to Optimize an Objective Function." <u>Journal of the</u>
- 546 American Statistical Association **58**(301): 236.

547

- Wenner, C. A. (1978). Anguillidae. In W. Fischer (ed.) FAO species identification sheets for
- 549 fishery purposes. W. A. F. A. 31). FAO, Rome.

550

- Wentworth, C. K. (1922). "A scale of grade and class terms for clastic sediments." Journal of
- 552 Geology **30**(5): 377-392.

553

- Whittaker, R. J., K. J. Willis and R. Field (2001). "Scale and species richness: towards a general,
- hierarchical theory of species diversity." Journal of Biogeography **28**(4): 453-470.

556

- 557 Wiggins, G. (2015). Larvae of the North American Caddisfly Genera (Trichoptera). University
- of Toronto Press, Scholarly Publishing Division.

559

- Williams, W. (2002). "Salinisation: A major threat to water resources in the arid and semi-arid
- regions of the world." Lakes & Reservoirs: Research & Management 4: 85-91.

562

- Woodward, G., D. M. Perkins and L. E. Brown (2010). "Climate change and freshwater
- ecosystems: impacts across multiple levels of organization." Philosophical Transactions of the
- 565 Royal Society B-Biological Sciences **365**(1549): 2093-2106.

566

- Wrona, F. J., T. D. Prowse, J. D. Reist, J. E. Hobbie, L. M. J. Levesque and W. F. Vincent
- 568 (2006). "Climate change effects on aquatic biota, ecosystem structure and function." Ambio
- 569 **35**(7): 359-369.

Figure 1

Regression analysis of diversity along a precipitation gradient.

Strongest relationships between environmental variables and (a-c) Annual Precipitation, (d-h) Fish Shannon Index, (i-j) Invertebrate Shannon Index, (k-l) Invertebrate Shannon Index. Fish diversity increases with increasing precipitation and Forested Riparian. Fish diversity decreases as NH4+ increases. Invertebrate diversity correlates negatively with Low Flow (LFPP). The second order polynomial fails to corroborate visual identification of a local maximum of macroinvertebrate diversity in the climate transition-zone. Conductivity, Surface Runoff, and Potential Evapotranspiration covary with precipitation and fish diversity. Slope, R^2, F-statistics, and *p*-values are reported in Appendix-regression.

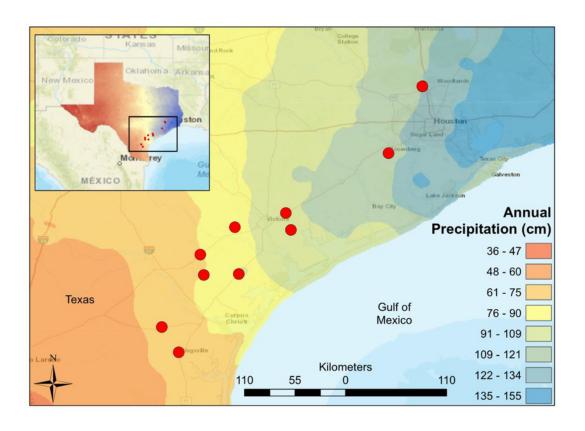




Figure 2

Map of sampling locations along the natural precipitation gradient in South-Central Texas.

10 USGS gaged Streams were sampled in the Spring of 2017. An annual precipitation overlay indicate that the sample sites span a gradient from 61 cm/yr in the Southwest to 121 cm/yr in the Northeast.

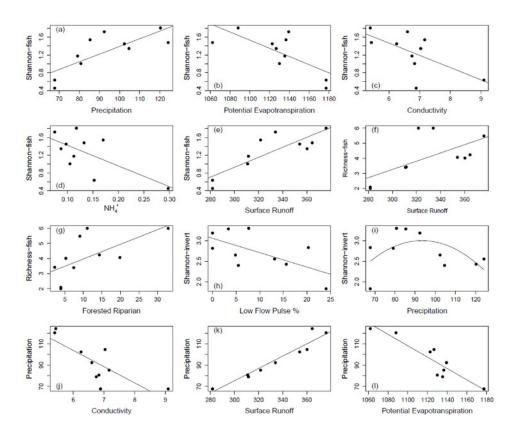




Figure 3

NMDS of (a) fish communities and (b) invertebrate communities.

Sites are grouped by precipitation regime; Semi-Arid (yellow), Mesic (green), and Sub-Humid (blue). Arrows indicate the strongest fitted environmental variables (p-value < 0.1) with length and direction corresponding to explanatory power. Semi-arid communities are dominated by livebearers and snails. Mesic and Sub-Humid communities contain a variety of sunfish, mayfly, caddisfly, true-bug, and beetle taxa. Sub-Humid communities contain different species of mayfly, caddisfly, true-bug, and beetle, but uniquely contain catfish, shrimp, and dragonfly taxa. Relative Humidity and Low Flow Pulse Percent correlates strongly with variation along the NMDS1, on which community grouping separate.

