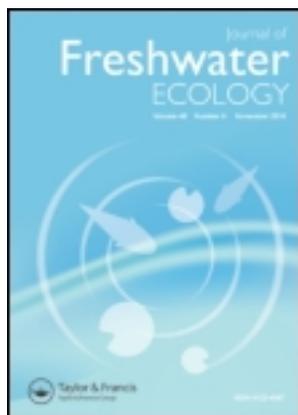


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EDITOR'S CHOICE ARTICLE

Effect of artificial light on the drift of aquatic insects in urban central Texas streams

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Light pollution can reduce night time drift of larval aquatic insects in urban streams by disrupting their circadian rhythms. Previous studies on larval insect drift show that disruption in drift leads to changes in reproduction as well as intraspecific and interspecific interactions. The purpose of this study was to conduct a preliminary investigation into the effects of extreme artificial light on insect drift in urbanized, high clarity spring systems of the karst Edwards Plateau, TX. We quantified taxa richness, diversity, and abundance in aquatic insect night time drift under two treatments (ambient night time light and artificial light addition) and among five streams using a paired design. Richness and diversity of drifting aquatic insects were similar between treatments but abundance was 37% less in the light addition treatment than that of the control. Effects of light addition on mean abundance was more notable in large streams with a 58% decrease in Simuliidae (compared to that of the control) and 51% decrease in Baetidae. Reduced drift from light addition suggests the potential of artificial lighting disrupting insect drift and consequently community structure. Results of this experiment support a growing body of knowledge on how urbanized systems influence stream communities.

Keywords: light pollution; stream ecology; urban ecology; drift; abiotic factors; Baetidae; Chironomidae

Introduction

Urbanization has the potential to cause numerous negative consequences for the physical and biological functioning of an aquatic ecosystem and looks to be a continuing ecological issue as the world's population migrates toward urban centers (Feminella & Walsh 2005). Physical processes of aquatic systems, including water quantity, stream morphology, and photoperiod, are susceptible to urbanization effects (Feminella & Walsh 2005). Photoperiod of aquatic systems can be disrupted by the presence of artificial night lights (i.e., street lights, safety lights, commercial lights) within and around the environment (Longcore & Rich 2004). Consequences of light pollution, through the increase in artificial lighting, are one frequently overlooked aspect of urbanization (Longcore & Rich 2004). In North America and Europe, 99% of the population is exposed to brighter than normal night skies with over 80% of the population experiencing skies brighter than that of a full moon (Navara & Nelson 2007).

To study the effects of light pollution on the environment, Longcore and Rich (2004) coined the term ecological light pollution that refers specifically to the disruption of ecosystem functioning by artificial night lights. Since the majority of organisms operate on a specific circadian rhythm, light pollution has the potential to impact the biological

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community by disrupting an organism's migratory and reproductive habits, thereby affecting biomass and population size through changes in interspecific and intraspecific interactions (Longcore & Rich 2004; Hölker 2010, Wolter, et al. 2010). Behavioral characteristics of larval aquatic insects are susceptible to ecological light pollution (Feminella & Walsh 2005). In particular, photoperiods of aquatic invertebrates are disrupted by artificial night lights located near or distant from aquatic ecosystems (Longcore & Rich 2004). Downstream drift is an essential part of dispersion among many aquatic invertebrates and is thought to assist in locating optimal living conditions (Müller 1974) or avoiding predation (Flecker 1992). As such, invertebrate drift rates peak immediately following dusk and continue through the night hours (Cloud & Stewart 1974; Ciborowski 1982; Brittain & Eikeland 1988). A number of families within orders Ephemeroptera, Diptera, Plecoptera, and Trichoptera undergo diel drift (Elliott & Minshall 1968; Pearson & Franklin 1968; Brusven 1970; Steine 1972; Cloud & Stewart 1974; Casey 1987).

Under laboratory conditions (Bishop 1969; Chaston 1969) and natural settings (Holt & Waters 1966; Perkin et al. 2011) aquatic invertebrate drift is reduced by artificial light. Disruption of night time drift from artificial lights causes insects to stop drifting over areas of suboptimum substrate or increased predation, leading to changes in recruitment for individual species and overall community composition (Blakely et al. 2006; Smith et al. 2009). Additionally, artificial light reduces the number of benthic invertebrates dispersing from the substrate, possibly increasing densities and intraspecific interactions among larvae (Palmer et al. 1996). Varying climates and ecosystem structures should be considered when studying drift to obtain a general picture of ecological significance (Brittain & Eikeland 1988) but, to date, effects of ecological light pollution are reported in streams at northern latitudes and researchers have yet to investigate the possible effects of artificial light in arid and semi-arid climates.

The purpose of this study was to conduct a preliminary assessment on the effects of additional artificial light on evening drift of larval insects in headwater streams within urbanized areas of the semi-arid Edwards Plateau region of central Texas. Urbanized streams within the Edwards Plateau provide a unique opportunity to assess the effects of artificial lights because of high water clarity (ranging from 0.1 to 10 Nephelometric Turbidity Units [NTUs]; Groeger et al. 1997), attributed to base flows predominantly from karst spring discharge (Hubbs 1995; Groeger et al. 1997; Saunders et al. 2001). With high water clarity, artificial lighting extends further into the water column. As such, we predicted that the light addition treatments would cause greater disruption of evening drift of aquatic insects than that reported at northern latitudes. The objective of this study was to quantify aspects of larval insect drift (diversity, abundance, and biomass) in urban streams under ambient night light conditions and under extreme light addition conditions.

Method

Study area

Study sites were located on two large spring runs with moderate levels of urbanization and three small spring runs with high levels of urbanization. The two large spring runs were San Marcos River (29.869395° N, -97.930194° W; Hays County, TX) and Comal River (29.710164° N, -98.129171° W; Comal County, TX). The three small spring runs were Onion Creek (30.188457° N, -97.71964° W; Travis County, TX), Bull Creek (30.40468° N, -97.789655° W; Travis County, TX), and Barton Creek (30.295928° N, -97.92642° W; Travis County, TX). Discharge at each site was near base flow

conditions. Each site included moderately flowing riffle or run mesohabitats with low levels of ambient light pollution. All streams had healthy riparian zones with moderate to high levels of riparian cover.

Physical habitat characteristics and water quality parameters were similar within large streams and within small streams. Large streams (>10 m width and >1.76 m³/s; San Marcos River and Comal River) were characterized by moderate depths (>0.5 m), swift current velocities (> 0.30 m/s), gravel to cobble substrates, and high water clarity (<10 NTUs). Smaller streams (<5 m and <0.12 m³/s; Barton Creek, Bull Creek, and Onion Creek) exhibited shallow depths (<0.3 m), slow current velocities (<0.1 m/sec), and sand to bedrock substrates. Water temperatures were 22–23 °C in large streams and cooler (<16 °C) in small streams. Dissolved oxygen levels were relatively low across all streams except at Bull Creek (10.6 mg/L). Specific conductance ranged between 491 and 650 μS/cm across all streams. Ambient light, measured directly above the water surface, was <2 lux across all streams.

Field collection

Five sites in Central Texas were sampled twice at night between November and December 2012 under two treatments: under ambient lighting (control) and with artificial light addition. Each night of sampling utilized one drift net (0.45 × 0.25 m, 500 μm mesh) and tested randomly one of the two treatments. Drift nets for each treatment were placed on separate evenings to ensure that the light addition treatment did not influence the results of the ambient light treatment. Additionally, only one location in each stream was tested at a time to minimize changes caused by environmental heterogeneity in large streams and due to minimal stream width and low water flow in small streams. Nets were set 30 minutes before sampling and captured drifting invertebrates for two-hour time periods starting within an hour after local sunset, because drift is highest during the first few hours after sunset (Cloud & Stewart 1974). Both ambient and light addition treatments were conducted within a week at each site to limit environmental differences between treatments.

Drift nets were placed in stream sections with the highest current velocities, typically downstream from a riffle, and supported by two metal fence posts. Nets were placed in the water column at least 5 cm above the substrate to minimize the likelihood of benthic insects crawling into the net. In large streams with a depth greater than the height of the net, the net was placed immediately below the water surface, since drift has been shown to be greatest in the surface current layer (Furukawa-Tanaka 1992). For the light addition treatment, two metal fence posts were set 1 m apart and supported a 1.2-m wooden plank with four portable work lights attached. Each portable work light held a 300-watt incandescent light bulb and was placed about 0.6 m above the stream surface. Lights were powered by a portable gas-powered generator. Artificial lights [mean = 1482 lx; standard error (SE) = 533] intensely illuminated a 1 × 4 m area and were placed 1 m upstream of the drift net. In large stream segments, the ambient light treatment included an additional two metal fence posts to simulate the setup of the light array. This addition was intended to control for the possibility of vegetation from the streams with high current velocity attaching to the metal posts and disrupting insect drift. After each sampling period, all contents collected from the drift nets were stored in 95% ethanol.

Laboratory analysis

Larval insects were separated from the rest of sample debris and placed in 95% ethanol for each sample site and treatment designation. Insects were identified to family level and

counted for abundance, richness, and diversity at each site and for each treatment. Biomass was estimated for the three most abundant taxa across all five streams. Lengths were measured for all individuals taken from small streams and for a subsample ($\leq 30\%$) of individuals taken from large streams (Allan 1982; McIntosh & Townsend 1996). Average biomass was estimated from average length using taxon-specific length–mass regressions (Table 2; Benke et al. 1999). Regression coefficients were unavailable for Leptohiphidae; therefore, coefficients for Ephemerellidae, a member of the same subfamily, were used instead. Total biomass was calculated by multiplying the mean family biomass per treatment by total number of individuals.

Data analysis

The experimental design was initially developed to accommodate a paired *t*-test in order to block differences in insect communities among streams. Drift densities of all taxa among five streams did not differ ($p > 0.05$) between treatments; however, trends in the data suggested an unanticipated stream size effect. The number of drifting taxa was more abundant and differences between treatments were more apparent in larger streams (San Marcos River and Comal River) than in smaller streams (Bull, Barton, and Onion Creeks). With low power ($n = 2$) to detect differences with parameter statistics for large or small streams, we calculated mean differences and 1 SE for dependent variables (density, richness, and total number) between the ambient light treatment and light addition treatment for all streams, and we visually estimated distributional differences among all orders and among most abundant families ($n \geq 82$ individuals). We considered treatment effects informative and detectable if mean differences and 1 SE were much less than zero (i.e., $\text{taxa richness}_{\text{light addition}} \ll \text{taxa richness}_{\text{ambient light}}$). Differences in biomass were tested with a paired *t*-test comparing results across all streams and within large streams.

Results

Among 10 collections, we captured 3190 individuals representing six insect orders [Ephemeroptera (53%), Diptera (24%), Coleoptera (11%), Odonata (6.6%), Trichoptera (4.7%), and Lepidoptera (1.3%)] and 36 families (Table 1). Abundant families were Leptohiphidae (33%), Baetidae (19%), Chironomidae (14%), Simuliidae (8.3%), Elmidae (6.8%), and Coenagrionidae (5.7%). The number of individuals per site was related to stream size with 90% of the insects captured from larger rivers (San Marcos River: 62%; Comal River: 28%). Overall, the number of drifting insects among all ambient light treatments ($n = 1979$) was greater than all light addition treatments (1212).

Mean differences (light addition – ambient light) plus 1 SE were $\ll 0$ for total number of larvae in all streams and large streams and for richness in large streams (Figure 1). Mean differences (light addition – ambient light) plus 1 SE among orders were $\ll 0$ for Diptera and Ephemeroptera in all streams and in large streams (Figure 2). Mean differences (light addition – ambient light) plus 1 SE among families were $\ll 0$ for Baetidae, Chironomidae, and Simuliidae among all streams and for Baetidae, Chironomidae, Coenagrionidae, Leptohiphidae, and Simuliidae among large streams (Figure 2).

Average biomass (mg) of drifting insects did not differ ($p > 0.05$) between treatments among the three most abundant families (Baetidae, Chironomidae, and Leptohiphidae) among all streams, large streams, or small streams (Table 2). However, total biomass, calculated by multiplying estimated average biomass by the total number of captured

Table 1. Total number of individuals (N), family richness (S), and Shannon–Weiner diversity (H') during the light addition and ambient light treatments at each of the five sampling sites.

| | Barton Creek | | Bull Creek | | Onion Creek | | Comal River | | San Marcos River | |
|-----------------------|--------------|----------|------------|----------|-------------|----------|-------------|----------|------------------|----------|
| | Light | No light | Light | No light | Light | No light | Light | No light | Light | No light |
| Coleoptera | | | | | | | | | | |
| Elmidae | 6 | 3 | 0 | 0 | 0 | 0 | 12 | 12 | 71 | 112 |
| Dropidae | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0 |
| Dystiscidae | 0 | 0 | 0 | 0 | 0 | 2 | 1 | 0 | 0 | 0 |
| Gyrinidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |
| Hydrophilidae | 0 | 0 | 2 | 1 | 1 | 3 | 0 | 0 | 0 | 0 |
| Lampyridae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 4 | 0 | 0 |
| Scirtidae | 0 | 0 | 1 | 0 | 26 | 48 | 1 | 2 | 17 | 10 |
| Tipulidae | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0 |
| Diptera | | | | | | | | | | |
| Ceratopogonidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 2 |
| Chironomidae | 11 | 1 | 5 | 12 | 7 | 22 | 73 | 134 | 78 | 115 |
| Culicidae | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0 |
| Empididae | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 5 | 2 | 0 |
| Ephydriidae | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0 |
| Simuliidae | 0 | 0 | 1 | 2 | 1 | 7 | 75 | 171 | 0 | 8 |
| Stratiomyidae | 0 | 0 | 2 | 1 | 0 | 1 | 0 | 1 | 6 | 3 |
| Tipulidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 |
| Ephemeroptera | | | | | | | | | | |
| Baetidae | 4 | 3 | 12 | 15 | 7 | 6 | 54 | 139 | 132 | 237 |
| Caenidae | 2 | 0 | 0 | 0 | 2 | 2 | 0 | 2 | 3 | 1 |
| Ephemeridae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 5 | 3 |
| Heptageniidae | 2 | 0 | 3 | 1 | 0 | 0 | 0 | 0 | 0 | 0 |
| Leptohyphidae | 22 | 0 | 9 | 7 | 0 | 0 | 41 | 68 | 377 | 512 |
| Leptophlebiidae | 1 | 0 | 1 | 0 | 0 | 0 | 4 | 0 | 5 | 18 |
| Lepidoptera | | | | | | | | | | |
| Crambidae | 0 | 0 | 0 | 0 | 0 | 2 | 7 | 12 | 9 | 12 |
| Odonata | | | | | | | | | | |
| Aeshnidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 2 | 2 | 3 |
| Calopterygidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 5 | 3 | 9 |
| Coenagrionidae | 9 | 0 | 11 | 12 | 0 | 0 | 6 | 20 | 44 | 80 |
| Cordulidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 2 |
| Libellulidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 |
| Protoneuridae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |
| Trichoptera | | | | | | | | | | |
| Brachycentridae | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 2 | 7 |
| Hydrobiosidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 2 |
| Helicopsychidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 |
| Hydropsychidae | 0 | 0 | 1 | 3 | 0 | 2 | 2 | 10 | 2 | 1 |
| Hydroptilidae | 0 | 0 | 1 | 3 | 0 | 0 | 5 | 4 | 1 | 5 |
| Philopotamidae | 0 | 0 | 0 | 2 | 0 | 1 | 3 | 6 | 26 | 44 |
| Polycentropodidae | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 6 | 7 |
| Total | 57 | 7 | 49 | 59 | 46 | 98 | 286 | 601 | 791 | 1196 |
| Shannon diversity | 1.706 | 1.004 | 2.053 | 1.988 | 1.365 | 1.656 | 1.89 | 1.918 | 1.769 | 1.842 |
| Family-level Richness | 8 | 3 | 11 | 11 | 5 | 9 | 15 | 21 | 21 | 25 |

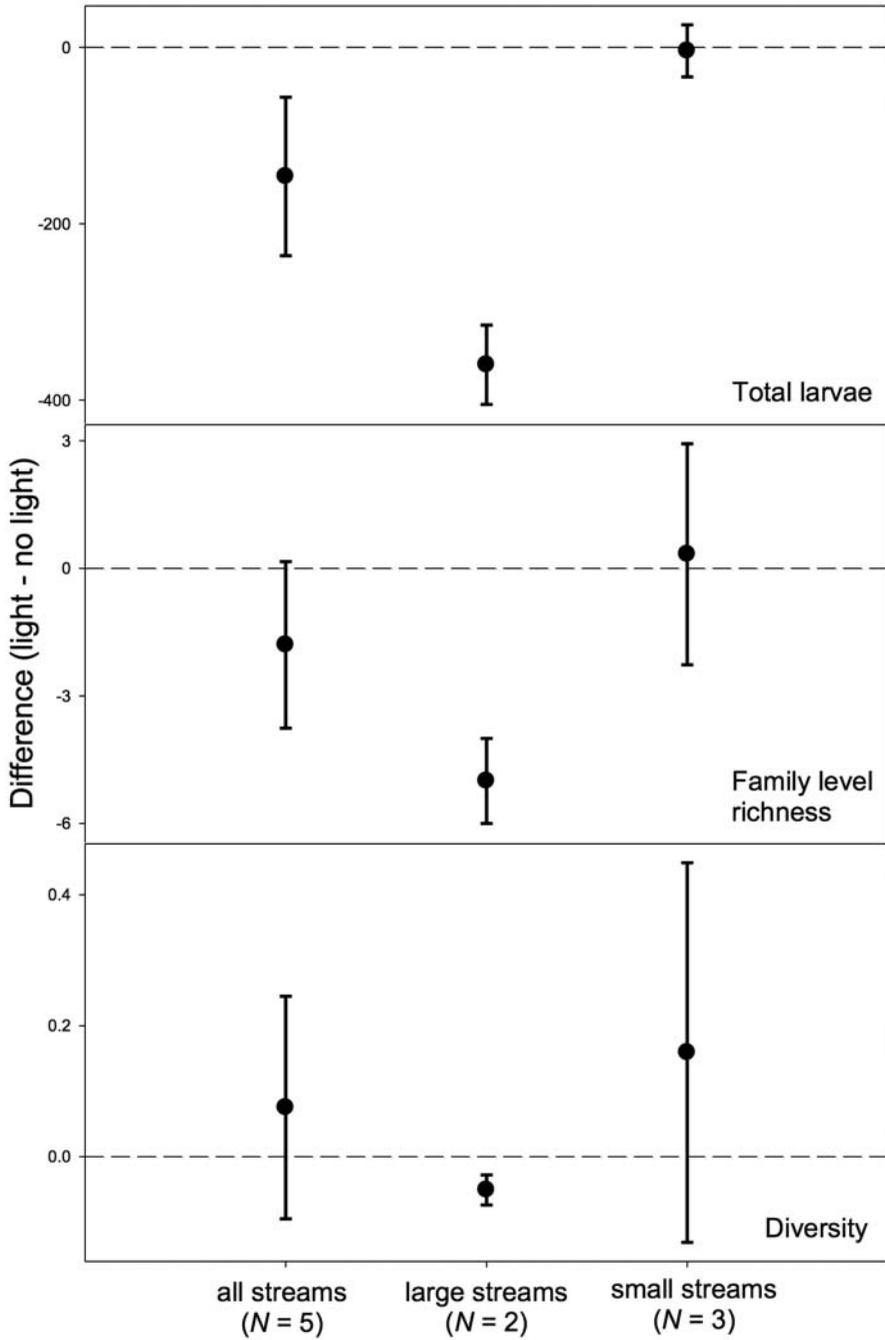


Figure 1. Mean abundance, family-level richness, and diversity \pm SE across all streams and within large and small streams from November 2012 to December 2012.

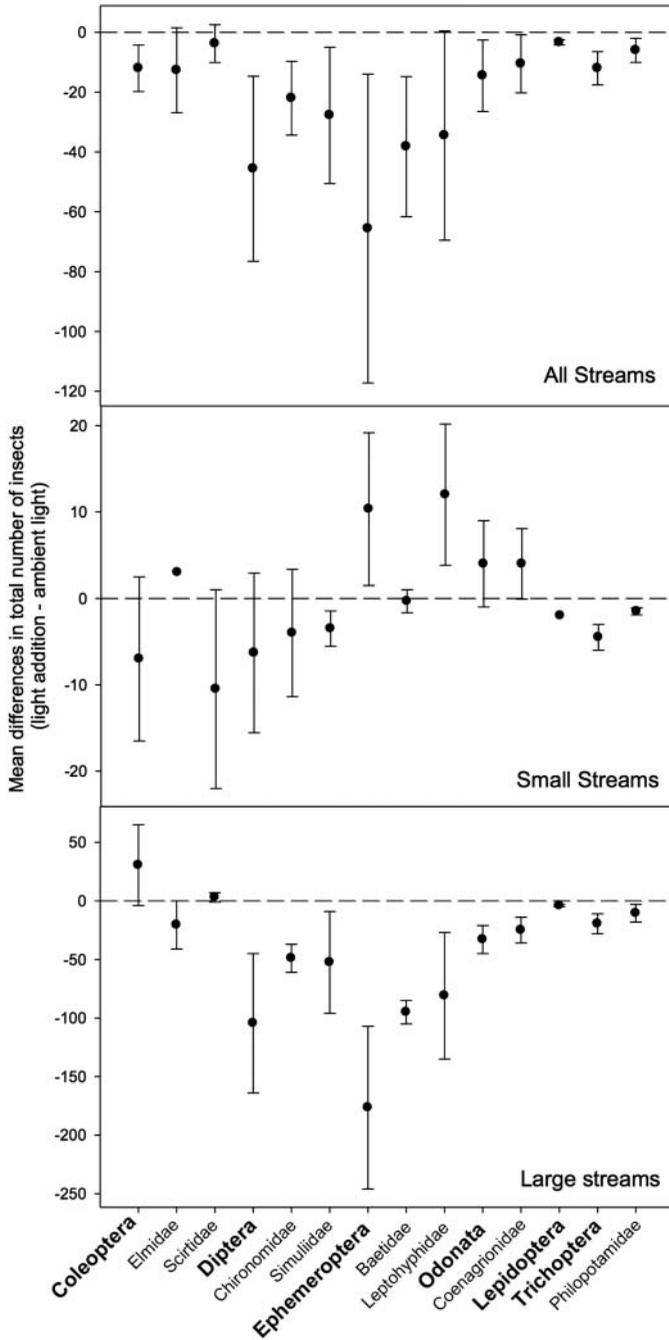


Figure 2. Mean abundance \pm SE for each order and family of aquatic insects across all streams and within large and small streams from November 2012 to December 2012. Difference for Lepidoptera in small streams and Elmidae in small streams based on one number.

Table 2. Average and total biomass (mg) for Baetidae, Chironomidae, and Leptohiphidae.

| | $\ln(M) = \ln(a) + b \times \ln(\text{body length})$ | | | | | | | |
|----------------------|--|----------|-------------|----------|-------------|----------|------------------|----------|
| | | | | | | | | |
| | | | | | <i>a</i> | | <i>b</i> | |
| | Bull Creek | | Onion Creek | | Comal River | | San Marcos River | |
| | Light | No light | Light | No light | Light | No light | Light | No light |
| <i>Baetidae</i> | | | | | | | | |
| Average biomass | 0.14 | 0.11 | 0.17 | 0.32 | 0.15 | 0.19 | 0.35 | 0.27 |
| Total biomass | 1.68 | 1.75 | 1.21 | 1.92 | 8.17 | 27.01 | 47.04 | 64.23 |
| <i>Chironomidae</i> | | | | | | | | |
| Average biomass | 0.03 | 0.05 | 0.12 | 0.07 | 0.06 | 0.06 | 0.05 | 0.06 |
| Total biomass | 0.14 | 0.57 | 0.85 | 1.54 | 4.26 | 7.84 | 3.73 | 6.41 |
| <i>Leptohiphidae</i> | | | | | | | | |
| Average biomass | 0.09 | 0.09 | NC | NC | 0.18 | 0.21 | 0.18 | 0.20 |
| Total biomass | 0.79 | 0.65 | NC | NC | 7.43 | 14.55 | 69.25 | 100.0 |

Note: NC = none collected.

individuals per treatment, differed for Chironomidae ($p < 0.01$) among all streams and Baetidae ($p < 0.01$) and Chironomidae ($p < 0.01$) within large streams (Figure 3).

Discussion

We found evidence to partially support our prediction that larval insect drift was less under additional light treatment than ambient light but only among a few taxonomic groups in larger streams. As such, insect drift within streams of high water clarity is not necessarily more susceptible to ecological light pollution but was disrupted similarly to drift reported outside of arid and semi-arid regions (Holt & Waters 1966; Perkin et al. 2011). Effects of additional artificial light were most evident for five families (Baetidae, Chironomidae, Coenagrionidae, Leptohiphidae, and Simuliidae) within larger streams. All are reported to undergo diurnal drift (Elliott & Minshall 1968; Pearson & Franklin 1968; Brusven 1970; Steine 1972; Cloud & Stewart 1974; Casey 1987), and our findings are consistent with reported decreases in drift abundance under artificial light for Baetidae and Simuliidae (Bishop 1969; Chaston 1969). Baetids decreased from 92% (number expressed as percentage of individuals drifting in that light level versus drifting in control with complete darkness) in a low light treatment (<0.2 lx) to 5.5% in the highest light intensity treatment (8.8 lx) and Simuliids decreased from 102% to 0% (Chaston 1969). Herein, we report a 48% decrease, on average, for Baetids and a 59% decrease for Simuliidae. Also, we report for the first time light effects on three of the families – Chironomidae, Coenagrionidae, and Leptohiphidae – that all experienced a decrease in drift due to the addition of artificial light.

The prediction of decreased drift due to the addition of artificial light was not supported within the smaller stream segments of this experiment. Small streams showed little difference in total abundance and richness. Each family captured in small streams on average showed no difference between the two treatments. A possible explanation for these unexpected results is that the smaller streams were part of a more urbanized

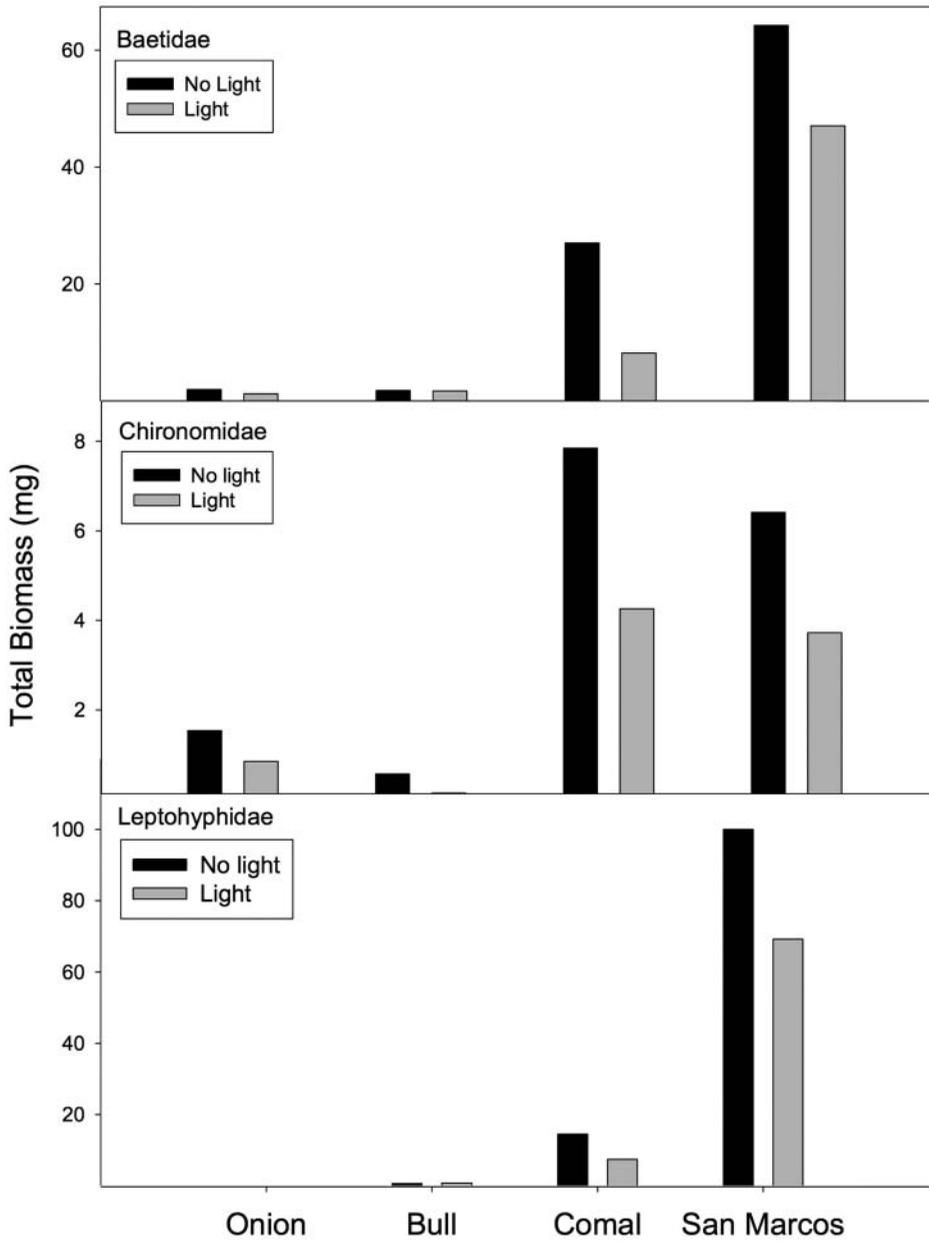


Figure 3. Total biomass (mg) for Baetidae, Chironomidae, and Leptohiphidae.

ecosystem, which can cause a simplification in the insect stream community (Morse et al. 2002; Gray 2004), potentially preselecting individuals with greater tolerance for urbanization and light tolerance. Taxa within orders Ephemeroptera, Plecoptera, and Trichoptera are considered pollution sensitive and have decreased in richness since 1996 in the lower reaches of Bull Creek (COA 2010). During this time, percentage of collectors has increased, which is often associated with streams of reduced water quality (COA 2010).

The decrease in drift shown across all streams and within larger streams has implications for community dynamics within streams and their adjacent riparian zones. Light pollution could cause insects to drop out early, disrupting drift and leading to changes in recruitment for individual species, and could simplify community composition as fewer insects make it downstream (Longcore & Rich 2004; Blakely et al. 2006; Smith et al. 2009). Since >80% of colonization movements are hypothesized to be caused by drift, any disruption due to light barriers could lead to an uneven distribution of insects within the benthic substrate (Townsend & Hildrew 1976). Urbanization, a major cause of the increase in light pollution, is also known to create subpar aquatic habitat, potentially causing drifting insects to land in areas of poor substrate (Blakely et al. 2006). Additionally, decreases in colonization due to light barriers could result in a loss of overall ecosystem productivity. Benthic invertebrates are often overlooked contributors to the aquatic ecosystem and play a key role in the aquatic trophic system (Covich et al. 1999). As stream insects are a large source of secondary productivity, an uneven distribution of drifting larval aquatic insects could result in a net loss of stream-wide productivity (Huryn & Wallace 2000).

Since many organisms operate on a circadian rhythm, ecological light pollution affects dispersal, communication, and predation among taxa, including aquatic insects, fish, terrestrial insects, bats, and migratory birds (Rydell 1992; Longcore & Rich 2004; Davies et al. 2012). Research in the area of ecological light pollution is still relatively new (Perkin et al. 2011) and there are many predictions to be tested. This experiment provides support for the hypothesis that extreme artificial light has negative effects on the abundance of drifting larval insects creating a need for conservation managers to look for ways to mitigate the effects. One possibility is the development of riparian buffer zones to decrease the amount of artificial light reaching the stream environment. Riparian buffer zones decrease the total amount and duration of light reaching a stream environment and influencing a sensitive biological process (Kiffney et al. 2004). Also, high-intensity lights have the greatest effect on insect drift (i.e., more than wavelength; Bishop 1969; Chaston 1969) and with light intensity varying by light source (Perkin et al. 2011), changes in the type of artificial lighting used (Hölker 2010, Moss, et al. 2010) could decrease the levels of light pollution. Installing low-intensity light bulbs, directing light away from the water's surface, or smart grids that turn off lights when not in use are among the possible solutions to light pollution, and already being implemented in major urban centers such as the city of Austin in central Texas. Tests on how stream communities respond to these higher efficiency lights could provide biological support to these initiatives as well as guide selection of future light efficiency measures.

This study provides specific evidence that artificial light disrupts the dispersal of larval aquatic insects in urban semi-arid streams; however, community-level implications of these changes are still to be determined. As an ideal biological indicator, benthic insects can be used to provide insight to environmental managers on the general health of an urban stream (Purcell et al. 2009). Given sufficient evidence that larval drift in urban semi-arid Texas streams decreases with light pollution, further studies could assess artificial light effects on larval insects with the addition of other urban stressors as urban streams can be difficult to restore with the numerous stressors at play (Purcell et al. 2009). However, it may be important to utilize larger and healthier stream environments in future studies, as treatment effects could be difficult to obtain within overly simplified communities of small urban streams. Future research will be necessary to determine how best to mitigate these effects and whether investments in mitigation have tangible environmental benefits.

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