

potentially implicates the cost of calcium uptake as a driver of increased metabolic rate under dilute conditions in organisms with calcified exoskeletons and suggests major phyletic differences in osmoregulatory physiology. Freshwater insects may be energetically challenged by higher salinities, while lower salinities may be more challenging for other freshwater taxa.

Freshwater ecosystems support a disproportionately large amount of the Earth's biodiversity [1–3]. Within these systems, salinity ranges naturally from practically deionized to ion rich [4] as a function of geology, rainfall, evaporation and hyporheic influences [5,6]. There are two dimensions of freshwater salinity change that appear to have significant ecological consequences. One dimension which appears to be predominant is the increasing salinization of freshwater, resulting from changing climate [7–11] and human activities such as hydraulic fracking [12] mountain-top coal mining [13], and irrigation of arid landscapes [14–16]. The second dimension involves systems becoming more dilute, due for instance to historic acid deposition [17–20]. It is apparent that both increasing and decreasing salinity regimes can have profound effects on biodiversity [13,21,21–25], but we do not yet understand how the physiological attributes of different freshwater species, and lineages, determine how salinity change affects performance at the organismal level.

Regardless of whether freshwater aquatic organisms originated from terrestrial ancestors (e.g., insects) [26–28] or invaded freshwaters from marine habitats (e.g., crustaceans) [29], they face the same physiological challenge of maintaining an internal environment (blood or hemolymph) that is strongly hypertonic to the surrounding medium. All freshwater animals must mitigate the inevitable effects of ion loss and water uptake [30–34], with some combination of regulating cuticular permeability and taking up ions against concentration gradients at the expense of ATP consumption [30,35,36]. To date, the energetic costs of osmoregulation in freshwater invertebrates remains poorly studied and major phyletic differences have yet to be revealed.

Aquatic insects typically thrive in relatively dilute conditions and often dominate fresh waters in terms of biomass, diversity and function [37–39]. Very few species of aquatic insects can survive in saline waters [31,40,41], and some entire lineages, such as mayflies, are being largely extirpated from anthropogenically salinized systems [13]. Aquatic insects are considered secondarily aquatic and derived from terrestrial ancestors [26,42] and it appears that actively regulating their internal salinity is a common attribute, even when physiologically stressed by increases in salinity [40,43–45]. In contrast, studies suggest that some freshwater non-insect taxa, many of which have marine ancestry, may be challenged by the same dilute conditions

that freshwater insects thrive in [46–48]. Thus, despite facing the shared physiological challenges that life in freshwater imposes, there are apparently fundamental species differences in osmoregulatory processes that remain poorly understood.

The literature provides contrasting information about the relationships between ionic concentrations and uptake rates. Organisms may have ion influx rates which are proportional to diffusive ionic loss rates, which are greatest when the concentration gradient is largest between hemolymph/blood and the surrounding water. For example, studies in rainbow trout show decreased calcium uptake in higher salinity waters [49]. However, freshwater insect ion uptake rates appear to uniformly increase with ion concentrations [36,40,50,51] (Cochran et al., in prep), with ion turnover rates potentially affecting the energy budgets of aquatic insects in waters with highly elevated major ions. This apparent energetic cost might explain developmental delays, reduced growth and other fitness consequences associated with higher salinities [44,52–55].

Existing data generally indicate that higher metabolic rates are associated with lower salinities in most freshwater species, implying a greater energetic cost of offsetting diffusive ionic losses in progressively dilute situations. This metabolic response to salinity has been seen in snails, gammarids, and other freshwater organisms with marine ancestry [56–59], but has not yet been linked to the transport of a specific ion. However, in aquatic insects we could only find a single paper exploring the linkage between metabolic rates and salinity in the salt tolerant mosquito, *Aedes aegypti* [60]. Curiously, Edwards reports that oxygen consumption is unaffected by environmental salinity. Given that we have ample evidence for a universal positive association between ionic concentration and uptake rates of calcium, sodium and sulfate in aquatic insects [36,40,50,51] (Cochran et al., in prep), we might expect that aquatic insect metabolic rates might increase with rising salinity to meet the increasing energetic demands of ion transport (as seen in African clawed frog, caridean shrimp, and others estuarine species [61–65]).

Here, we asked if fundamentally different physiological responses to salinity exist between aquatic insects and other aquatic species. We explored relationships between salinity, ion transport rates and oxygen consumption in several species (including three aquatic insect species and three other aquatic species- two scuds, *Hyalella azteca* and *Gammarus pulex*, and one snail, *Elimia sp.*). These species were specifically chosen to compare aquatic insects (with terrestrial ancestry) to species of non-insects (with marine ancestry) known to have a metabolic response to salinity. Specifically, we used respirometry to measure metabolic rates in each species in response to waters with different conductivities. Further, we assessed whether

movement of specific ions may be more energetically costly by performing ion flux experiments using radiotracers in waters with different conductivities and ion compositions.

Methods

Animal rearing and collection

Neocloeon triangulifer (mayfly) was reared using previously described methods [36,66]. Briefly, *N. triangulifer* (WCC-2 clone) was originally isolated for culture from White Clay Creek (WCC), Chester County, PA, USA and rearing methods were developed at the Stroud Water Research Center (SWRC; Avondale, PA) [67]. At North Carolina State University, cultures were maintained in 4-qt glass Pyrex dishes, lined with WCC periphyton plates) [67] and aerated to maintain oxygen saturation. Naiads were reared at 21–23 °C, 14 h:10 h light:dark photoperiod from egg hatch until they were relatively mature (approximately 3 weeks old) and approximately equal in size.

Hyalella azteca (scud) were purchased from Goliad Farms (Goliad, TX), packaged with ice packs, and shipped overnight to NC State University. *Gammarus pulex* were ordered from Critters Direct (New Bern, NC). After being received, all organisms were acclimated in ASW at room temperature with dried leaves for food.

Elimia sp. (snail) were collected from Beaver Creek in Greensboro, North Carolina (35.999°, -79.712°) by removing specimens from rocks with forceps. Live snails were packaged with ice packs and transported to NC State University for experimental use. Snails were acclimated in Artificial Soft Water (ASW) at room temperature. The caddisfly, *Hydropsyche betteni* was collected from a riffle flowing out of Yates Mill Pond in Raleigh, North Carolina (35.718°, -78.686°) by removing larvae from rock bottoms with forceps. Animals were transferred back to NC State University in a mason jar with mesh substrate and acclimated in ASW at room temperature. *Drunella cornutella* were collected from the Basin Creek in Hays, North Carolina (36.375°, -81.144°) using a D-frame kick net in riffles and animals were transferred back to NC State University in an aerated cooler. All field collected species were acclimated in ASW at room temperature.

A dendrogram (fig. 1) was assembled using phyloT (<https://phylot.biobyte.de/>), which automatically generated the tree based on the NCBI taxonomy database and visualized in iTOL (<https://itol.embl.de/>). Branch lengths are arbitrary and only represent taxonomic structure.

Water preparation

CrystalSea Marine Mix (MM) (Marine Enterprises International, Baltimore, MD, USA) was added to reverse-osmosis deionized water to create the desired experimental concentration (see Table 1). Waters were monitored for 24h to ensure desired final conductivity.

For waters where conductivity was held constant and Ca content was manipulated, we diluted the 136MM water with DI water (either 85% or 75% 136MM by volume) and titrated CaCl_2 until the conductivity of the water reached $136\mu\text{s cm}^{-1}$ and remained there for 24h.

Artificial soft water (ASW) is our routine culture media and was prepared as previously described [36,60].

Respirometry

We used an 8-channel, fiber-optic, intermittent flow respirometry system (Loligo Systems, Tjele, Denmark) with AutoResp (version 2.1.0; Loligo Systems, Tjele, Denmark) software to obtain standard metabolic rate (SMR) measurements. Chambers (1.3 ± 0.1 mL) were fitted with a glass spacer ring and stainless-steel mesh, which kept organisms separated from the continuously stirring magnet glass stir bar. Each chamber and corresponding fiber-optic channel recorded measurements of oxygen consumption for one individual. Chambers were submerged in an aerated, temperature-controlled water bath (approx. 4 L). Water bath temperature was controlled using a programmable heater/chiller (12108-30; Cole Palmer, Vernon Hills, Illinois), which cycled water between the respirometry bath, and a heat exchange coil submerged in the heater/chiller reservoir for the entire experiment to maintain a temperature of 20°C . For each experimental run (in which 8 individuals were evaluated), background oxygen consumption measurements ($n = 3-4$ 15-min cycles) were first taken from blank respirometry chambers using the intermittent respirometry setting in AutoResp 2.0. A Gilson Minipuls peristaltic pump (Gilson Inc., Middleton, WI) refreshed the chamber medium with surrounding bath water (ASW) at the initiation of each 15-minute cycle (300 s flush, 350 s wait, 250 s measure). Then, an individual was introduced to each of 8 chambers and intermittent flow respirometry was performed for 2 to 3 cycles (30–45 min). Individuals underwent no prior acclimatization or fasting.

After 2 to 3 cycles the AutoResp software was changed to the “closed” setting and chambers were no longer refreshed by the peristaltic pump until the oxygen level in the chambers dipped to around $80\%\text{O}_2$. Previous experiments have suggested that oxygen saturations above $80\%\text{O}_2$ appear to not be stressful in most species as they are well above most species critical oxygen

tension (P_{crit}) at 20°C [68]. When the oxygen levels in the chambers dipped to around 80%O₂, the peristaltic pump was turned back on to refresh the chambers back to 100%O₂. After at least one hour, the source water for 4 of the 8 chambers was changed to a different treatment water (see table 1). The same protocol was repeated with half of the chambers maintained in ASW and half exposed to the treatment water (oxygen level dipping to 80%, then being refreshed to 100%) for at least two hours. This was repeated twice for each treatment (yielding n=16 for ASW for each species and n=8 for other treatments).

AutoResp software reports M.O₂ in $\mu\text{g g}^{-1} \text{h}^{-1}$, and accounts for differences in O₂ capacity of water due to salinity (when different salinity is specified in the software). A one-way ANOVA and Tukey's multiple comparisons test found no significant differences between runs for a species, so all runs were combined for future analyses. We used the (q0.2) approach [69], which uses quantiles to place SMR above the lowest 20% of observations for each individual before kPa fell below 10.5 (which corresponded with the relatively flat portion of each run for each individual).

To analyze effects of salinity, data was plotted using GraphPad Prism (v6, GraphPad Software, La Jolla, CA, USA). Differences among salinities were assessed using a one-way ANOVA and Tukey's multiple comparisons test. All salinities were compared to every other salinity. Values are presented as mean \pm SEM unless otherwise indicated.

Ion flux experiments

Individuals were 19 days old for all ²²Na and ⁴⁵Ca uptake experiments. Radioactive experimental waters were made with each treatment water, spiked with ⁴⁵CaCl₂ or ²²NaCl (PerkinElmer, Billerica, MA, USA). Exposure activities ranged from 130 to 220 Bq mL⁻¹. Exposures were measured with PerkinElmer Wallac Wizard 1480 Automatic Gamma Counter (Shelton, CT, USA) or Beckman LS6500 Multipurpose Scintillation Counter (Beckman Coulter, Brea, CA, USA) immediately before the experiments began.

100mL high-density polyethylene beakers with 15mL of radioactive exposure water were gently aerated, sealed with ParaFilm™ and spatially randomized. Each beaker held 3 individuals, 1 per time point (3, 6, and 9-hours for dual-labeled with ²²NaCl and 12, 24, and 36-hours for ⁴⁵CaCl₂) and there were 8 replicate beakers. At each time point, individuals were removed from the radioactive exposure waters by gently pipetting them into a mesh strainer (collecting any residual radioactive water in a waste container) and gently blotting dry. The organisms were then rinsed in two consecutive water baths of the corresponding unlabeled exposure water to

remove loosely adsorbed ions from the exoskeleton. For ^{45}Ca experiments, individuals were also rinsed with 0.05 M EDTA and 0.1 M L-ascorbic acid sodium salt due to the adsorptive nature of Ca on exoskeletons [70]. After rinsing, individuals were blotted dry, weighed, and digested in 500 μL of Soluene 350 (PerkinElmer) in a 20 mL glass vial at 28°C. After 48h, they were neutralized with 500 μL of glacial acetic acid and 12 mL of scintillation cocktail (PerkinElmer Ultima Gold uLLT).

The uptake of each ion was plotted against time and uptake rates were calculated as the slopes of linear regression analysis using GraphPad Prism. Mass-specific calculations were based on wet weights. We applied appropriate corrections for spill-over and quench, and only measurements with lumex values <5% and error rates <10% were used in analyses. Flux rates were compared among groups using a one-way ANOVA with Tukey's multiple comparisons test using GraphPad Prism.

Results

3.1 Respirometry

There was an apparent decrease in metabolic rates (M.O_2) with increased salinity in *G. pulex* (fig. 2A), *H. azteca* (fig. 2B) and *Elimia sp.* (fig. 2C). For example, *G. pulex* had a strong negative correlation between water conductivity and M.O_2 $r(5)=-0.97$, $p=0.14$, however the relationship wasn't statistically significant. Metabolic rates were unaffected by changes in salinity in *D. cornutella* (fig. 2D), *N. triangulifer* (fig. 2E) and *H. betteni* (fig. 2F).

Ion flux experiments

There was an apparent increase in sodium uptake rates with increased salinity for *G. pulex*, *H. azteca* and *Elimia sp.* (fig. 3). We found a strong positive relationship between sodium uptake rates and conductivity for *G. pulex* ($r(3)=0.96$, $p=0.18$), *H. azteca* ($r(3)=0.98$, $p=0.11$) and *Elimia sp.* ($r(3)=0.96$, $p=0.16$), however the relationship was not statistically significant.

In Marine Mix waters, calcium uptake rates decreased with increased salinity for *G. pulex* (fig. 4A), *H. azteca* and *Elimia sp.* (table S1). Oxygen consumption rates were also positively correlated with calcium uptake in *G. pulex* ($r(5)=-0.93$, $p=0.02$) (fig. 4B), *H. azteca* ($r(3)=-0.99$, $p=0.03$) and *Elimia sp.* ($r(3)=0.91$, $p=0.26$) (table S1).

In *G. pulex*, when water conductivity was held constant, but calcium content was manipulated, calcium uptake rates were positively correlated with Ca concentration ($r(4)=0.97$, $p=0.02$) (fig. 5A). Further, calcium uptake was positively correlated with metabolic rate ($r(4)=0.94$, $p=0.05$) (fig. 5B).

Discussion

Existing literature indicates lower salinities are associated with increased metabolic rates in gammarids, snails, common carp and cutthroat trout [56–59,71–74] (fig. 6A). These findings suggest a greater energetic cost of offsetting diffusive ionic losses in progressively dilute situations in these species. However, ecological [13,21] and organismal/physiological [45,70,75,76] studies indicate more saline conditions are more stressful to most aquatic insects, even when given the opportunity to acclimate [40,66]. Thus, despite the common need for all freshwater organisms to regulate their internal environments in hypotonic surroundings, there are clearly physiological differences among lineages that are yet to be described.

Therefore, we first explored relationships between salinity and oxygen consumption rates in six species, three insects (from terrestrial ancestry) and three non-insects (from marine ancestry), across a gradient of salinity. Commensurate with previous studies [56–59], we found that *G. pulex*, *H. azteca* and *Elimia* sp. had decreased metabolic rates with increasing salinity. Though not studied here, studies have shown some estuarine and brackish water species demonstrate increased metabolic rate with increasing salinity (potentially to meet the increasing energetic demands of ion transport) [61–65,77] (fig. 6B). However, this is not the case for all euryhaline species, for example there is evidence for decreased oxygen consumption with increased salinity in the euryhaline tilapia (*Oreochromis mossambicus*) [78]. Here, we show for the first time that aquatic insects do not show a metabolic response to salinity. We therefore propose a third potential scenario of energetic impact of osmoregulation in hypotonic environments (fig. 6C). In this scenario, organisms may have increased ion fluxes with increasing salinity; however, unlike the scenario in fig. 6B, metabolic rate may not change with salinity. Here, it appears that aquatic insects primarily rely on shifting their energetic stores to meet the increased demands of ion transport, rather than altering their metabolic rate to cope with increased salinity, leading to delays in growth and development [55,76]. To our knowledge, a similar metabolic response to salinity has only been seen anecdotally in aquatic insects, in the salinity tolerant mosquito *Aedes aegypti* [60], the Chinese mitten crab [79], the European eel [80], Brine shrimp [81] and the giant freshwater prawn [82].

Next, we asked if the changes in metabolic rates observed with increased water conductivity in the three non-insect species could be linked to the transport rates of a particular ion. We found that Na uptake rates were positively correlated with Na concentration and associated conductivity (fig. 3). This is consistent with previous work in aquatic insects, where a positive association between Na uptake and the Na concentration of the water is the rule (Cochran et al., in prep) [36,40,51,75]. However, Na uptake rates were negatively correlated with metabolic rate (fig. 3), which suggests that Na uptake does not explain the increased energetic cost for these species under dilute conditions. Alternatively, we found that Ca uptake was negatively associated with conductivity for all three non-insect species in these experimental waters. This finding differs strongly from all the insect studies we have done where we have always observed a positive association between Ca uptake and Ca concentration (Cochran et al., in prep). In *Daphnia*, acclimation to dilute water has been shown to increase Ca uptake [83]. However, our findings are consistent with other studies in rainbow trout (*Salmo gairdneri* Richardson, 1836) and tilapia (*Oreochromis mossambicus* W. K. H. Peters, 1852) where increased salinity was associated with decreased calcium uptake [63,65].

One interpretation of our data is that under dilute challenge, non-insect species (or those with calciferous exoskeletons) prioritize the increased uptake of Ca despite its apparent energetic cost (e.g., increased Ca uptake is stimulated by dilute challenge). An alternative, though not mutually exclusive possibility is that other ions (e.g., Na, SO₄) have a suppressive impact on Ca uptake rates when those ionic concentrations increase. An example of this type of suppression was observed by Scheibener, who showed that Na is antagonistic to SO₄ transport in *N. triangulifer* [45].

We provide two lines of evidence that link calcium uptake rates to metabolic rates in non-insects. First, we demonstrate a stimulation of calcium transport under increasingly diluted conditions that was linked to increased metabolic rates. This may be associated with energetic costs of increasing calcium transport to offset diffusive losses. Second, when we held total conductivity constant (136 μ S/cm) and manipulated calcium concentration from 1.7 to 17.3 mg/L, we see a positive association between calcium uptake rates and metabolic rates. This finding suggests that calcium transport in these animals is largely linked to ATP consumption. However, we do not observe this pattern in aquatic insects. Why are they so different?

One possibility is that the rates of ion transport do not change in response to salinity. We reject this notion, based on several studies which show that aquatic insect transport rates appear to always be concentration dependent. For example, Scheibener et al. 2016 found that

sodium transport was positively associated with sodium concentration in *H. betteni* [75]. Scheibener et al. 2017 found that sulfate uptake was positively associated with sulfate concentration for *N. triangulifer*, *Drunella sp.*, *Ephemerella sp.*, *Isonychia sp.*, and *Maccaffertium sp.* [45]. A recent study by Cochran et al. has found a positive association between calcium, sodium and sulfate uptake and calcium, sodium and sulfate concentration (respectively) in 8 species of aquatic insect species (including mayflies, caddisflies and mosquitoes) (Cochran et al., in prep). While these studies show that this is the case in naïve animals (e.g., animals exposed to short term or abrupt salinity changes), we also show that animals will acclimate to high salinity conditions by reducing uptake rates towards control rates, though their uptake rates are still concentration dependent. For example, Orr et al. shows that calcium, sodium and sulfate uptake rates are positively associated with calcium, sodium and sulfate uptake rates (respectively) in *N. triangulifer* after chronic exposure to elevated ion treatments [51]. Similar trends have also been seen in *Callibaetis floridanus* after chronic exposure to a wide range of salinity treatments [40]. We also see acclimatory change in *N. triangulifer* exposed to progressively more dilute matrices with increases in transport rates observed in acclimated animals that more closely approximate standard rearing conditions than do higher salinity acclimated animals [3.6].

A second possibility is that aquatic insects maintain somewhat constitutive expression of ion transporting ATPases and utilize secondary transporters to modulate overall uptake rates when salinity changes. However, proteomic data from the gills of *N. triangulifer* exposed chronically to different salinities show that V-ATPase, Na/K ATPase and Ca ATPase change [84]. *N. triangulifer* and *Callibaetis coloradensis* have shown increases in ionocytes (mitochondria-rich osmoregulatory structures) with more dilute rearing condition [36,85]. Edwards also show increases in mitochondria in the anal papillae of *A. aegypti* with dilute rearing conditions [60]. More studies are needed to further address the function and response of these transporters.

A third possibility is that aquatic insects shift the allocation of their energetic resources to osmoregulation at the expense of growth and development [55,76]. We are still confronted with the conundrum of the energetic linkages between osmoregulation and metabolism. One perspective is that the most energetically favorable situation for aquatic organisms is to have blood that is isotonic to the external media [86]. This perspective is clearly not the case in insects where they appear to thrive in dilute environments and suffer when the external environment approaches isotonicity with the internal media [21,43]. It is tempting to speculate

that the terrestrial derived aquatic insect osmoregulation system is optimized to very large transmembrane potentials between apical cells and the external media but to our knowledge this has never been measured.

Intriguingly, we generally see very low rates of Ca uptake in aquatic insects relative to Na and SO₄ and often must use longer exposure durations in our Ca uptake studies. It appears that relatively slow Ca uptake is a common feature of aquatic insects who have chitinous exoskeleton, (as opposed to calciferous in crustaceans) [47,66]. In a comparative Ca study radiotracer with Hydropsychidae and Ephemerellidae, Poteat [70] found Ca uptake rates were so slow that it was necessary to lower stable Ca exposure concentrations to facilitate uptake of ⁴⁵Ca. Further, Orr [51] provided histological evidence for possible Ca damage to Malpighian tubules under Ca rich conditions, suggesting that excessive Ca uptake is potentially toxic in aquatic insects. In contrast, Ca is a significant component of crustacean exoskeletons and molluscan shells [87–90], and those species appear to increase their metabolic rates to support Ca uptake under dilute conditions.

It remains unclear to what extent these phyletic differences are linked to the evolutionary origins; however, we posit that the evolutionary history of these species may have some impact on their physiological response to salinity. Freshwater systems are comprised of organisms from many different lineages. Phylogenomic analysis of aquatic insects show that there is no clear common ancestor, but rather that multiple invasions of water from land that occurred over millions of years [26–28]. Alternatively, 30 clades of crustaceans, such as the scuds *G. pulex* and *H. azteca*, have invaded freshwater from marine origins [29]. Likewise, all living freshwater mollusks also evolved from marine origins (including 15 gastropod clades) [29].

Here, we posit that there are three distinctive relationships among salinity and metabolic rates in aquatic organisms (fig. 6). In the first, organisms (such as gammarids) have decreased metabolic rates with increasing salinity [56–59]. In the second, metabolic rate increases with increased salinity (as seen in estuarine species) [61–65]. In the third, we show for the first time that aquatic insects have no metabolic response to increased salinity. We also provide evidence that in non-insect species with a metabolic response to salinity, it appears as though Ca is the driver rather than Na. Altogether, we suggest that both the decreased demand for Ca and lack of metabolic response to salinity in aquatic insects could provide a physiological explanation for aquatic insects thriving in dilute environments, relative to other taxa. However, future work in non-insect aquatic lineages lacking a marine ancestry and/or a calciferous exoskeleton (or shell)

would be informative in determining whether that relationship is a function of the biology of aquatic insects or ancestry.

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Competing interests

No competing interests declared.

Data availability

Data available on request from the authors.

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Figures and Table

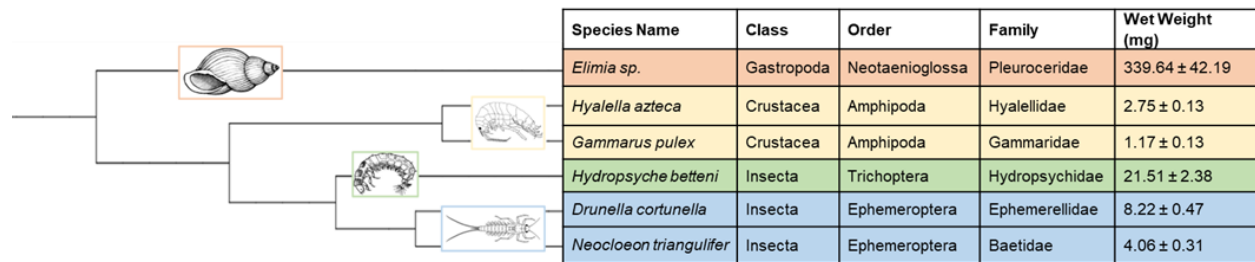


Fig. 1. Dendrogram showing taxonomic structure of species used in these experiments (generated using phyloT software) and species information (including class, order, family and wet weight (mg)).

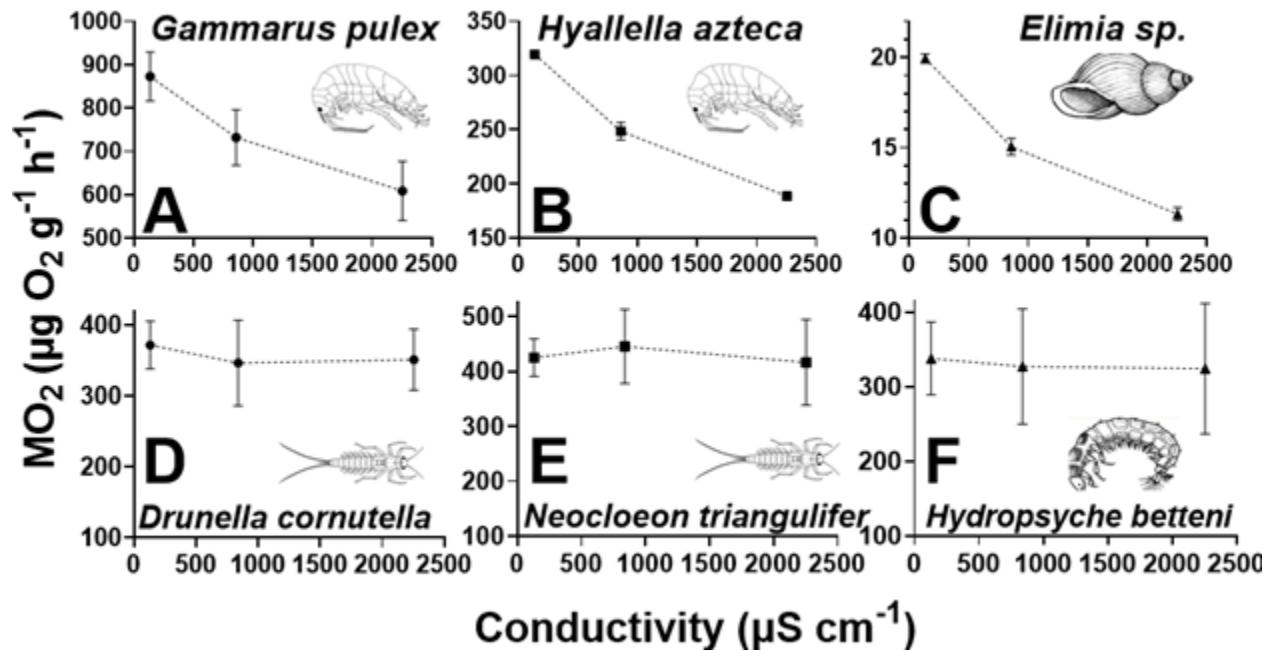


Fig. 2. The influence of salinity on oxygen consumption rates (M.O₂) in aquatic invertebrates. Lineage specific patterns emerge with crustaceans (A,B) and a snail (C) showing increased consumption with decreasing salinity and mayflies (D,E) and a caddisfly (F) showing no change (n=16 for ASW; n=8 for other treatments).

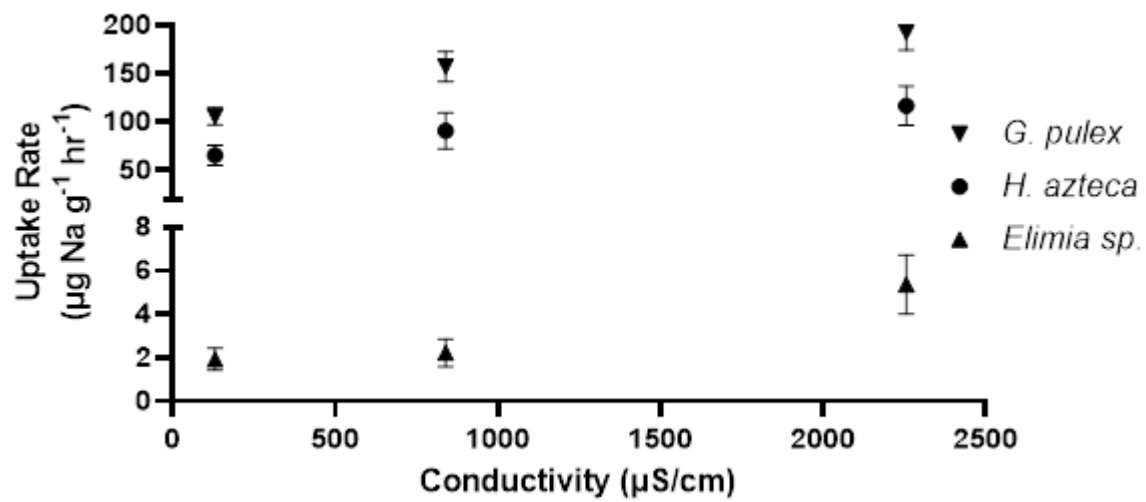


Fig. 3. Sodium uptake rates are positively associated with salinity and sodium concentration in *G. pulex*, *H. azteca* and *Elimia sp.* when exposed to a serial dilution. (mean±SEM) (n=8)

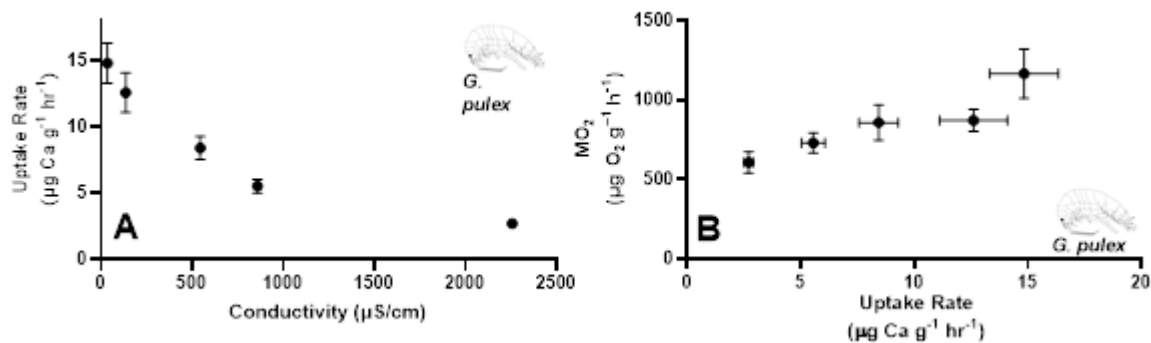


Fig. 4. Calcium uptake rates are negatively associated with salinity and calcium concentration in *G. pulex* when exposed to a serial dilution (A). Calcium uptake rates were positively associated with oxygen consumption rates (B). The same pattern is observed in *H. azteca*. (n=8)

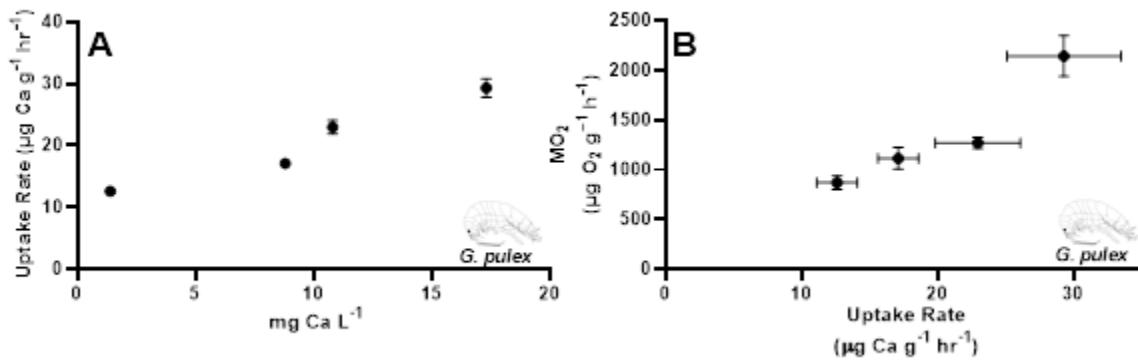


Fig. 5. Calcium uptake rates are positively associated with (A) calcium concentration and (B) oxygen consumption rates in *G. pulex* when conductivity is held constant, but calcium varies from 1.7 to 17.3 mg Ca L⁻¹. (mean±SEM.) (n=8)

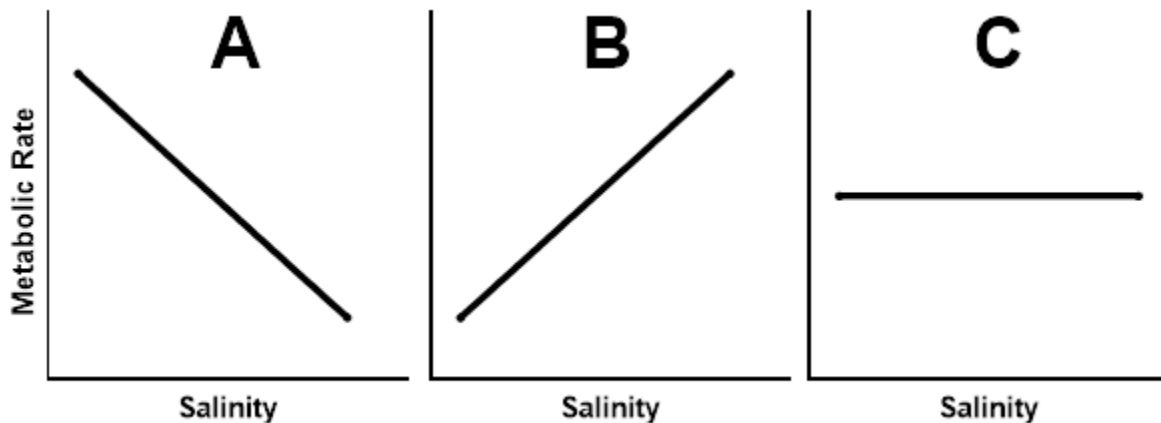


Fig. 6. Three models of energetics and fluxes in aquatic organisms. **A.** Metabolic rates are higher in dilute conditions and associated with the increased need to offset diffusive losses (e.g., snails). **B.** Metabolic rates increase with salinity to meet increasing the energetic demands of ion transport (e.g., observed primarily in some marine/euryhaline species). **C.** Metabolic rates do not change with salinity (e.g., aquatic insects).

Table 1. Water chemistry for all experimental waters. Conductivity is reported in $\mu\text{S cm}^{-1}$. Ions and TDS are all reported in mg L^{-1} . *S represents total S (as SO_4). All waters were sampled, filtered, and verified by NC State University's Environmental and Agriculture Testing Services Lab (ICP-MS).

Name	Conductivity	g MM L^{-1}	Na	Ca	S*	Mg
35MM	35	0.02	9.7	0.3	0.9	0.9
136MM	136	0.077	31.6	1.4	3.5	3.7
ASW	136	-	15.5	10.8	7.3	3.1
85%136 + Ca	136	0.065 (+30.6 $\text{mg CaCl}_2 \text{ L}^{-1}$)	27.1	8.8	2.9	3.1
75%136 + Ca	136	0.057 (+34.9 $\text{mg CaCl}_2 \text{ L}^{-1}$)	23.8	17.3	2.6	2.7
560MM	560	0.486	131.1	4.4	15.4	14.0
853MM	853	0.778	201.7	6.9	23.	21.6
2256MM	2256	2.094	339.5	11.4	32.3	42.7