Physiological capacity of Cancer setosus larvae — Adaptation to El Niño Southern Oscillation conditions

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ABSTRACT

Temperature changes during ENSO challenge the fauna of the Pacific South American coast. In many ectotherm benthic species pelagic larvae are the most important dispersal stage, which may, however, be particularly vulnerable to such environmental stress. Thermal limitation in aquatic ectotherms is hypothesized to be reflected first in the aerobic scope of an animal. Here we present results on whole animal oxygen consumption and on the activities of two metabolic key enzymes, citrate synthase (CS) and pyruvate kinase (PK) of Cancer setosus zoal larvae, acclimated to different temperatures. Larvae acclimated to cooler temperatures (12 and 16 °C) were able to compensate for the temperature effect as reflected in elevated mass specific respiration rates (MSR) and enzyme activities. In contrast, warm acclimated larvae (20 and 22 °C) seem to have reached their upper thermal limits, which is reflected in MSR decoupling from temperature and low Q10 values (Zoea I: 1.4; Zoea III: 1.02). Thermal deactivation of CS in vitro occurred close to habitat temperature (between 20 and 24 °C), indicating instability of the enzyme close to in vivo thermal limits. The capacity of anaerobic metabolism, reflected by PK, was not influenced by temperature, but increased with instar, reflecting behavioral changes in larval life style. Functioning of the metabolic key enzyme CS was identified to be one possible key for larval limitation in temperature tolerance.

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1. Introduction

Temperature is often referred to as the main factor determining biogeographical distribution of marine organisms. The physiological background of temperature dependent latitudinal ranges of marine ectotherms has been subject of numerous studies and during the last decades our knowledge on temperature limitation and acclimation processes has expanded. However, most of these studies refer to adult animals. More recently, attention focused on early life stages, which are generally hypothesized to be more vulnerable to stress conditions than juveniles or young adults (Anger, 2001). Early life stages are important for species distribution and recruitment especially in benthic coastal ectotherms (Cowen and Sponaugle, 2009).

Temperature affects all levels of biological organization ranging from cellular to organism level (Cuderley and St-Pierre, 2002). A mismatch between the oxygen demand of tissues and the supply by the circulatory and ventilatory system is the first mechanism restricting survival at unfavorable temperatures (Heilmayer et al., 2004; Pörtner, 2001; Pörtner, 2004; Storch et al., 2009). The thermal tolerance thresholds (pejus and critical limits) in the concept of oxygen and capacity limited thermal tolerance (OCLT) presented by Pörtner et al. (2005) describe the shortage of aerobic capacity due to unfavorable temperatures. These thermal limits can be shifted by different (acclimation) mechanisms, e.g. changes in the kinetic characteristics of enzymes (Heilmayer et al., 2004; Sokolova and Pörtner, 2001).

Two temperature sensitive key enzymes regulating energy provision in the form of ATP were chosen as proxies of thermal acclimation and limitation at an enzymatic level: citrate synthase (CS) as an indicator of animal aerobic metabolism (Hochachka et al., 1970) and pyruvate kinase (PK), which represents the potential for glycolytic flux (Childress and Somero, 1979; Johnston et al., 1977; Lemos et al., 2003). Investigations on metabolic enzymes in larval stages are scarce. Studies in shrimp larvae show that CS activity is inversely correlated with growth during ontogeny (Lemos et al., 2003). Metabolic capacity thus influences crucial performances and fitness at ecosystem level.

The model species, Cancer setosus (synonymous Cancer polyodon Poeppig 1836), studied herein covers a wide latitudinal cline of about 44° (Fischer and Thatje, 2008; Garth and Stephenson, 1966). The commercial value of this species for the Chilean and Peruvian artisanal fishery increased during the last decades (Sernapesca, 2006; Thatje et al., 2008; Wolff and Soto, 1992). The El Niño Southern Oscillation (ENSO), with drastically changing water temperatures between La Niña and El Niño, strongly affects the abundances and distribution range of this commercially important crab. Increasing temperatures during El Niño
events are discussed to be one of the main factors causing mass mortalities of this species (Arntz et al., 1988). The early ontogeny, which in C. setosus consists of five planktotrophic zoeal stages (Zoea I–V = Z I–V) and one megalopa before reaching the first crab stage (Quintana and Saelzer, 1986), is considered as the most delicate part within the life cycle of brachyuran and in particular cancrid crabs (Anger, 2001; Weiss et al., 2009a). High nutrient-content in eggs of C. setosus at high latitudes, supports their plasticity to respond to latitudinal and seasonal changes in temperature (Fischer et al., 2009). The relationship between environmental temperature and metabolic acclimation has been discussed in a large variety of studies, but knowledge of temperature effects on the physiology of crab larvae, especially concerning the ontogenetic development, is scarce. Furthermore, it remains speculative how larval instars are capable to compensate for the drastic temperature changes as caused by ENSO.

The present paper examines the influence of temperature on metabolism (oxygen consumption and enzyme activities) of larval C. setosus from the Antofagasta region in order to evaluate their capacity to respond to fluctuations in temperature as encountered by larvae during ENSO.

2. Materials and methods

2.1. Sampling and maintenance of adults

Ovigerous C. setosus (carapace width: female A = 12.8 cm CW, female G = 14.0 cm CW) were caught in February 2007 by fishermen of the “Caleta Colosso” (23°45′ S, 70°27′ W) by scuba diving and immediately transferred to the laboratory of the Instituto de Investigaciones Oceanológicas of the Universidad de Antofagasta, Chile. Animals were maintained individually in flow-through seawater (12 l) at ambient temperature ~16.0 °C and salinity 34 psu during ENSO.

2.2. Experimental set-up

Freshly hatched larvae were collected in filters receiving water from the overflow of the aquaria. Since most larvae hatched at night, samples were taken every morning. Filters were cleaned every evening to ensure daily larval age did not vary by more than 12 h (after Lovrich et al., 2003). Solely actively moving larvae were transferred to 100 ml bowls with 16 °C filtered seawater and afterwards were transferred to the corresponding experimental chamber. Acclimation temperatures were chosen to simulate LN conditions (12 °C), normal conditions (16 °C) slight EN (20 °C) and EN conditions (22 °C). For each acclimation temperature an initial number of 2000 larvae was cultured (maximum 10 individuals per bowl), water was changed daily and larvae were checked for molts or mortality and fed ad libitum with living Perumytilus purpuratus.

2.3. Oxygen consumption

Randomly selected larvae of the same age from the midst of Zoea I (ZI), Zoea III (ZIII) and Zoea V (ZV) at each acclimation temperature were chosen (as available — see Table 1) for measurements of the acute temperature dependence of oxygen consumption rates. The middle of each instar period was determined by using the larval development periods described in Weiss et al., 2009a. 8 Zoea I instar larvae, 3 Zoea III larvae, and 2 Zoea V larvae per replicate were transferred to an acclimation bath, where they were allowed to acclimate to the measurement temperature (12, 16, 20 or 22 °C) for 20 min to avoid shock reactions and to exclude weak larvae (those which died during the acclimation period) from further analyses. Afterwards larvae were gently transferred into 1.0 ml glass caps with fully aerated seawater and sealed with silicon membrane lids. Care was taken that no air bubbles were enclosed. Oxygen micro-optodes (needle-type, fiber-optic microsensor, flat broken tip, 140 μm) were inserted through the silicon membranes into the glass caps. Each measurement consisted of 3 replicates and 1 blank (glass cap filled with aerated seawater). Blanks were run in order to correct for bacterial oxygen consumption. Each combination of temperature and larval instar

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Table 1

Overview about conducted experiments. ZI, ZIII, ZV — Zoeal instars, P — postmolt, A — premolt, “+” — experiment conducted, “−” — experiment not conducted due to insufficient larval material.
was tested 3 times (see Table 1). The caps and the optodes were fixed in a mounting rack, which was placed into the temperature controlled basin. Optodes were connected to a 4-channel microsensor oxygen meter (PreSens GmbH, Regensburg, Germany), and oxygen was continuously recorded once every 15 s until depletion occurred by a minimum of 5% oxygen. During the experiments the experimental setup was gently shaken once every 5 min to avoid the development of oxygen gradients within the glass caps. Prior to experiments, optodes were calibrated with aerated seawater (see above) for 100% oxygen, and with a saturated sodium dithionite (Na₂S₂O₄) solution for 0% oxygen. Experimental temperatures were kept constant (±0.5 °C) in temperature controlled water baths.

2.4. Enzyme assays

Samples for the determinations of enzyme activities of citrate synthase (CS) and pyruvate kinase (PK) were taken in the post- (P) and premolt (A) period of the Zoea I (ZIP, ZIA), Zoea III (ZIIIPI, ZIIIA) and Zoea V (ZVP, ZVA) instar of larvae reared at the four acclimation temperatures (12, 16, 20, 22 °C) and analyzed following a modified method of Sidell et al. (1987) (CS) and a modified method of Simpfendörfer et al. (1995) (PK), which were adopted for small sample size and measurement in a microplate reader. Three replicates were collected for each “instar × acclimation temperature” combination as available (see Table 1).

Frozen samples were homogenized in extraction buffer (75 mM Tris–HCl, 1 mM EDTA; pH 7.6) at a ratio of ~0.3 μl per 1 μg larval DW (dry weight) to get a 1:3 (w/v) ratio with a Branson Sonifier 450 (0 °C, output control 8, duty cycle 50%, 15 min). Cell debris was removed by centrifugation for 5 min at 7400 g and 0 °C with an Eppendorf centrifuge 5410R.

2.4.1. Protein content

The concentration of soluble protein in the extracts was measured according to Bradford (Bradford, 1976). The samples were diluted 1:5 with 0.9% NaCl before being applied in duplicate (5 μl) on microplates. Subsequently 250 μl dye reagent (Biorad protein assay 5000006, diluted 1:5 with aqua dest) were added and the optical density was measured at 620 nm in a microplate reader (FLUostar Galaxy, BMG). Bovine serum albumin (BSA, 0–3.5 μg per well) was run in parallel as standard.

2.4.2. Citrate synthase

Citrate synthase (CS) (E.C. 4.1.3.7) is a key regulatory enzyme in the tricarboxylic acid (TCA) cycle and was chosen as an indicator of aerobic capacity. Due to the limited volume of extract the assay according to Sidell et al., 1987 was adapted for use in a microplate reader and under controlled (below ambient) temperature conditions.

Homogenates (2 μl/well) were assayed in 200 μl of 75 mM Tris–HCl buffer (pH 8.0), 0.25 mM DTNB (5,5′-dithio-bis-(2-nitrobenzoic acid)) and 0.4 mM acetyl-CoA. 0.4 mM oxalacetate was added to start the reaction (omitted for the blanks). The microplates were incubated at the respective temperature (12, 16, 20, 22, 28 °C) on a thermostated aluminum block; and the coupled oxidation of NADH was measured quickly every 5 min for 30 min (6 measurements) in a microplate reader at 405 nm and room temperature after which time the plate was returned to the aluminum block for further incubation. Enzyme activity in units per mg protein (U gprot⁻¹) was calculated using a standard curve produced from NADH (0.5 mM) and corresponding to 7 to 35 μM NADH groups.

2.8. Statistical analyses

All data were tested with the Jackknife distance test (Kezdi et al., 2002) to exclude outliers from analysis. The effect of acclimation temperature Tᵐ, measurement temperature Tᵐ (Kelvin) and body mass DW (μg dry mass) on metabolic activity (oxygen consumption, CS activity, PK activity) was assessed by means of analysis of covariance (ANCOVA) according to the model

\[ \text{In(rate)} = a + b₁/Tᵐ + b₂ + \ln(DW) + D₁ + D₂/Tᵐ \]

where variables D₁ and D₂ attain values specific for Tᵐ (12, 16, 20, 22 °C). Only the significant terms of each model are presented in the Results chapter. The relationship between ln(CS activity) and the inverse of Tᵐ was not linear over the whole temperature range, i.e. this model could not be applied (see Results). Instead, we introduced Tᵐ as a categorial variable to test for effects of W and of Tₙo.

3. Results

3.1. Oxygen consumption

Larval oxygen consumption is predicted from measurement temperature (Tᵐ), acclimation temperature (Tₐ) and larval body mass (DW) by the model

\[ \text{ln(Oxy)} = 8.8781 - 4207.9393 \times 1/Tᵐ - 0.0885 \times \ln(DW) + D₁ + D₂/Tᵐ - 0.0034 \]

where D₁ = 0.0678, -0.0953; 0.0153; 0.0122 and D₂ = -2154.3805; -1001.2125; 1102.7057; 2052.8873 for Tₐ = 12, 16, 20, 22 °C.

N = 256, F = 15.112, R² = 0.33, and p < 0.0001 for the whole model and each term. Please note that 1/Tᵐ is adjusted to mean = zero in the interaction term in order to make the test for the main effects independent of the test for interaction (“centered polynomials”).

Note that the model (Fig. 1) predicts larval oxygen consumption for just any combination of larval body mass and measurement temperature; whereas our experiments indicate that the “body mass × temperature” space where larvae perform aerobically is limited (see Discussion). Outside the pejus range larval oxygen consumption rates will most probably level off.

In general whole organism oxygen consumption of C. setosus larvae increased with progressing development. However, depending on acclimation temperatures the increase varied without any discernible pattern. Mass specific respiration rates (MSR) showed significant differences between acclimation temperatures (p = 0.040, F = 2.81, df = 3), measurement temperature (p = 0.0001, F = 84.60, df = 1), body mass (p = 0.0002, F = 14.38, df = 1) and in the interaction term of and “acclimation temperature × measurement temperature” (p = 0.0086,
As expected, MSR of larvae increase with measurement temperature. Thermal sensitivity of oxygen turnover (expressed as Q10) decreased with increasing acclimation temperature from 2.3 at 12 °C acclimated Zoea I larvae to 1.4 in 22 °C acclimated larvae (data not shown). This decrease is even more pronounced in ZIII larvae, showing a Q10 of 2.28 at an acclimation temperature of 16 °C and a Q10 of 1.02 at 22 °C acclimation temperature.

3.3. Citrate synthase

The citrate synthase (CS) activity of *C. setosus* larvae differed significantly depending on acclimation temperature (p = 0.0003, F = 6.34, df = 3), assay temperature (p < 0.0001, F = 21.20, df = 1) and body mass (p = 0.0007, F = 11.56, df = 1). CS activity was found to be highest at an acclimation temperature of 12 °C, and remained virtually unchanged at the higher acclimation temperatures. This decrease is even more pronounced in ZIII larvae, showing a Q10 of 2.10 at an acclimation temperature of 16 °C and a Q10 of 1.02 at 22 °C acclimation temperature.

3.2. Citrate synthase

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The CS activity decreases with increasing body size. The number of assay temperatures did not allow exact calculations of Arrhenius breakpoint temperatures (ABT) with acclimation temperature and instar. Nevertheless, the ABT of CS seems to be located between 20 and 24 °C, as seen in the Arrhenius plot (Fig. 2).

3.3. Pyruvate kinase

Larval PK activity is predicted best from measurement temperature (Tm) and larval body mass (DW, μg) by the model

$$\ln(\text{PK}) = 11.8344 - 4158.7783 \times \frac{1}{T_m} + 0.1469 \times \ln(DW)$$

N = 844, F = 548.09, R² = 0.57, and p < 0.0001 for both slopes and for the whole model. PK activity showed a continuous increase with measurement temperature (Fig. 3). The Q10 values were lower for freshly hatched larvae (ZIP, Q10: 1.17) than for ZI larvae in the premolt phase (ZIA: 1.83 ± 0.14), but in ZII and Z IV Q10 values are higher in the postmolt phase than during the premolt phase (ZIIPIIP, Q10: 1.86 ± 0.097; ZIII, Q10: 1.34 ± 0.722; ZVP, Q10: 1.41; ZVA, Q10: 1.39). PK activity increased continuously with body mass, with slightly higher values in the beginning of the ZIIIA and ZVA instar than at the end of the corresponding instar.

The CS/PK ratio was determined for 16 and 20 °C, temperatures which supported complete zoeal development. Here the CS/PK ratio was not dependent on acclimation temperature, but values decreased significantly from Zoea I to Zoea V (p < 0.0001; N = 65; df = 5; F = 8.65).

4. Discussion

Temperature limitations of *C. setosus* larvae were found to be reflected in the functioning and activity of the metabolic key enzyme CS. Our results indicate that larvae undergo cold acclimation associated with an activity increase of CS at an acclimation temperature of 12 °C, and remained virtually unchanged at the higher acclimation temperatures. CS activity increased with increasing assay temperature, until about 20 to 22 °C, but was significantly lower at the assay temperature of 28 °C (Fig. 2) which is also reflected in different Q10 values for the lower (12–20 °C, Q10: 1.5 ± 0.36) and upper temperature range (20–28 °C, Q10: 0.45 ± 0.15), the latter indicating thermal inactivation of the enzyme.

![Fig. 1. Model of oxygen consumption of larval *Cancer setosus* throughout their development. Points show real data.](image1)

![Fig. 2. CS activity in units per mg protein (U gprt⁻¹) at different assay temperatures (12–28 °C) of Zoea I *Cancer setosus* larvae. Arrhenius equation (black line) only fits for assay temperatures 12–20 °C, indicating thermal inactivation of the enzyme at higher temperatures. Arrhenius equation: \( \ln(\text{CS}) = 6.57 - 3035.05 \times \frac{1}{T_m} \).](image2)
Standard metabolic rate of an organism typically increases exponentially with increasing temperature between critical temperatures, which border the wider thermal tolerance window including the passive range until anaerobic metabolism sets in (Pörtner et al., 2005). As Zoea larvae are actively swimming in the water column, larval oxygen consumption comprises standard metabolism and oxygen demand for swimming, and can be best described as routine metabolism. Although an exponential pattern of routine metabolism couldn’t be observed in larvae of the congener Cancer irroratus (Sastry, 1979) and other decapod larvae such as Taliepus dentatus (Storch et al., 2009), swimming Cancer setosus larvae obviously display an exponential increase of their routine metabolism with increasing temperature (see Fig. 1). The window between critical limits of swimming larvae would thus result narrower than in the resting larvae.

The developed model (Fig. 1) allows us to predict the MSR of Cancer setosus larvae for any combination of body mass and measurement temperature, but experiments indicate a limitation of the aerobic performance.

As expected, mass specific oxygen consumption decreases with body mass. Interestingly the slope of this decrease is much lower (−0.08) than expected theoretically and as we know from empirical data (−0.25). An almost linear increase of individual respiration rates with DW normally is only found during periods of increased growth (Heilmayer et al., 2004; Hoegh-Guldberg and Manahan, 1995). This finding is also reflected in the extremely high cumulative growth rates of Cancer setosus larvae (Weiss et al., 2009a) when compared to other decapod crustacean larvae (Anger, 1995). When acclimated to higher temperatures (20 and 22 °C) larval routine oxygen consumption failed to increase with measurement temperature, demonstrated in low Q10 values at higher acclimation temperatures (see Results). Part of this loss in thermal response may be due to reduced metabolism. This indicates that aerobic larval metabolism of warm acclimated animals is reaching its upper limit and has no or only very limited capacities to adjust to further temperature increments. As stated by Hoegh-Guldberg and Pearse (1995) a lack of temperature compensation indicates that larval development occurs close to the upper possible temperature limit. In cold acclimated larvae a higher thermal plasticity was found, indicated by higher Q10 values well within the thermal range. Zoea III acclimated at 12 °C show particularly high oxygen consumption rates and corresponding high levels of CS activity.

An elevated CS activity at cold temperatures indicates metabolic cold compensation (Lemos et al., 2003; Sokolova and Pörtner, 2001). CS activity measured at different assay temperatures increased with increasing temperature before declining in vitro at temperatures warmer than 20 °C (Fig. 2), indicating onset of thermal deactivation of the enzyme. Such deactivation is manifest in reduced activities in all groups at 28 °C assay temperature even when compared to 12 °C assay temperature. This means that the temperature optimum for the functioning of CS lies close to the optimum temperature for larval development and growth (~20 °C) (Weiss et al., 2009a) in the Antofagasta region. Although denaturation temperatures are usually found far beyond the naturally experienced temperatures of (temperate and cold-water) ectotherms (Sokolova and Pörtner, 2001) as supported by the present PK data, a similar discontinuity in CS activity has been described for Littorina saxatilis, where the deactivation temperature of the enzyme in vitro lies equally close to the high ambient temperatures encountered by the animal (Sokolova and Pörtner, 2001). This suggests that the failure of CS at high temperatures may substantially contribute to the thermal limitation of larvae.

The functioning of CS is restricted in all larval instars. Our results also indicate that the ABTs of CS in higher instars of warm acclimated larvae are shifted to warmer temperatures. Further investigations with even more assay temperatures are needed to substantiate this finding. Such a shift in ABT with acclimation temperature may suggest that higher instars show a certain capability for warm acclimation and are therefore less vulnerable to elevated temperatures. This may correspond with the local oceanographic conditions in the experimental region, where younger larvae most likely drift into the Antofagasta bay and are retained within a cyclonic current at elevated temperatures for longer periods of time. According to the present data an acclimation of the CS properties occurs under the local conditions in the experimental area. This may support elevated instar dependent temperature optima of growth rates in Cancer setosus Zoea II + III instars (Weiss et al., 2009a).

In light of the temperature changes associated with ENSO Zoea I instars display very limited tolerance as indicated by the lack of thermal compensation in respiration rates and the low denaturation temperature of CS, (sensu Sokolova and Pörtner, 2001) The ABT of CS between 20 and 24 °C (Fig. 2) (Table 5) indicates thermal inactivation of CS closely preceding the acute lethal limit of Cancer setosus Zoea I larvae, which lies between >24 and ~30 °C (Weiss et al., 2009b and preliminary experiments). Thus, the dysfunction of mitochondrial enzymes and more generally a loss in aerobic capacity may contribute to mortality under acute heat stress, as during EN events, especially in warm water regions like the Antofagasta bight.

The moderate warm acclimation capacity of the larvae matches the slight seasonal temperature variations of ~4 °C (SHOA, 2009) in Antofagasta, more than the larger (up to 10 °C) temperature fluctuations associated with EN events. Oxygen consumption measurements and recent studies of elemental composition and phenotypic plasticity of body size and spine characteristics (Weiss et al., 2009a; Weiss et al., 2009b) indicate that Cancer setosus larvae already reach their thermal limits at ~22 °C.

CS activity is changing in vivo depending on the instar and is clearly declining towards the Zoea V (Table 5). A decline in CS activity through ontogenetic development can be related to the 450-fold increase in body size (Weiss et al., 2009a) and thus allometric effects from instar to instar. Another reason for this decrease may lie in the reduced requirement for locomotory activity and therefore aerobic capacity after transition from planktonic to benthic life style which occurs within the megalopa (Lemos et al., 2003).

In contrast to CS activities, pyruvate kinase (PK) activities were not affected by acclimation temperature indicating that PK is not contributing to thermal compensation or displays sufficient capacity at any acclimation temperature. PK activity increased continuously with assay temperature from 12 to 28 °C, with no signs of thermal inactivation (Fig. 2). PK activity increased with increasing larval size, and thus the CS/PK ratios decreased with increasing instar, as would be expected from allometric relationships. Higher glycolytic TCA (tricarboxylic acid) capacities in later larval instars indicate higher capacities for anaerobic metabolism. These findings correspond with larval life styles, as they improve their capacity to hunt prey and escape from predators during their ontogeny (Lemos et al., 2003). Furthermore, higher PK over CS activities at higher temperatures indicate a strategy to stabilize energy metabolism by use of...
anaerobic capacities which help to survive short-term thermal extremes. In higher instars (ZIII and ZV) PK activity is more pronounced in the beginning of the instar than in the end (also reflected by Q10 values higher in the beginning of an instar than in the end), which may indicate low PK activity during ecdysis. During the premolt phase food uptake is stopped, which obviously weakens the larva, as visible in the CHN values and the high mortality during ecdysis (Weiss et al., 2009a). Low PK activity during the energy consuming molt shows that larvae can only revert to a limited anaerobic capacity, which in total may contribute to the high mortality of larvae during ecdysis.

5. Conclusion

C. setosus zoeal instars show high temperature sensitivity in aerobic metabolism. Respiratory and CS capacities show compensation in the cold but are limited at warmer temperatures. Anaerobic capacities display no compensation, but a higher anaerobic than aerobic capacity may be associated with slowing larval life style. Later in ontogeny when the capacity to forage by hunting improves, PK activity rises indicating higher capacities for anaerobic metabolism. The results of this study indicate that C. setosus larvae are able to display a certain thermal compensation in the cold, but larvae obviously already live at their upper tolerance limits. Limitation of aerobic metabolic pathways seems to be responsible for a restricted thermal tolerance of C. setosus larvae during EN events.

Acknowledgments

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References