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Effects of temperature and salinity on survival and growth of the amphipod Hyale crassicornis (Gammaridea, Hyalidae)

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Abstract

This study explored the survival and growth of the marine amphipod *Hyale crassicornis* under different conditions of ambient temperature and salinity in the laboratory. In 96-h exposure experiments at a salinity of 30, H. crassicornis tolerated temperatures between 6 and 28° C ($> 90\%$ survival). The 96-h LT_{50} value (median lethal temperature) at high temperature range for the amphipod was 32°C. The values at low temperature range for adults and juveniles were 3.2 and 4.2° C, respectively, indicating that adults have a higher tolerance to low temperature than juveniles. At 25° C, the amphipods tolerated salinities between 10 and 50 (with $>80\%$ survival in 96 h). The 96-h LS₅₀ values at high and low salinity ranges were 56 and 2, respectively. Growth of the juveniles from instars 1 to 6 was determined under different combinations of temperature and salinity. Growth rate of juveniles cultured at 25° C and a salinity of 20 was the highest $(0.114 \text{ mm day}^{-1})$ among the combinations of 15-25°C and 10-40 salinity. The moult increment of body length was affected by both temperature and salinity. Yet the intermoult duration of the amphipods was affected by temperature only but not by salinity.

Keywords: Amphipod, Hyale crassicornis, Gammaridea, Hyalidae, temperature, salinity, growth, survival

Introduction

Environmental variables exert a significant impact on the physiology of all organisms. Temperature and salinity are generally regarded as the two major environmental determinants affecting the survival and growth of marine invertebrates (Kinne 1970, 1971). For instance, Brown and Bert (1993) showed that temporal variations in temperature and salinity limited the survival of juvenile stone crabs Menippe adina Williams and Felder, 1986. Both survival and growth of the penaeid shrimp, *Penaeus* chinensis (Osbeck, 1765), were affected by salinity and temperature (Chen et al. 1995, 1996).

Biological knowledge of environmental tolerance and growth is essential to the development of optimal culture conditions for animals (Hardy et al. 1994). Many marine invertebrates have specific temperature or salinity requirements at different developmental

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stages (Kinne 1970, 1971; Thomas and Rice 1992; Charmantier and Charmantier-Daures 1994; Guerin and Stickle 1997a). Understanding of the physiological requirements and tolerance limits in different growth phases of animals to environmental conditions will be helpful in further studies on their life history and population distributions. Such knowledge is also essential to the development of sensible pollution indices for the evaluation of environmental deterioration (Rosas and Ramirez 1993; Alcaraz et al. 1997). For instance, Alcaraz et al. (1997) reported that post-larvae of the white shrimp Penaeus setiferus (Linnaeus, 1767) had a lower tolerance to thermal stress when the animals were exposed to nitrogenous contaminants such as ammonia and nitrite.

The amphipod Hyale crassicornis (Haswell, 1880) (Gammaridea, Talitroidea, Hyalidae) is distributed in intertidal areas of Kiama, New South Wales, Cape Naturaliste and Western Australia (Barnard 1974). This species can also be found beneath rocks and pebbles, often associated with seaweed in the intertidal zone of Tolo Harbour, Hong Kong (own observation). Yet, information on the distribution of the species in South-East Asia is limited. Few studies on hyalid amphipods in Hong Kong waters have been undertaken (Moore 1986). Jiang and Zhou (1982) and Moore (1986) recorded four species of the genus *Hyale* found in Hong Kong but none of them was identified as H. crassicornis.

Amphipods have been suggested to be good biological indicators for the detection of early environmental degradation (Zanders and Rojas 1992). Interest has arisen in developing H. crassicornis as a pollution indicator organism in Hong Kong waters but biological knowledge of this species is very limited. This study aims to: (1) investigate the survival of the amphipod at different temperatures and salinities, and (2) elucidate the relationships between its growth and these two environmental variables. Such information would enhance our understanding of the biology of this amphipod and assist the development of pollution bioassays.

Material and methods

Survival at different temperatures and salinities

Acclimation. Amphipods were collected from Tolo Harbour, Hong Kong and maintained in three 3000-litre outdoor culture tanks at the Marine Science Laboratory of The Chinese University of Hong Kong. The tanks contained natural sea water with a turnover rate of \sim 2600 l day⁻¹. The animals were chosen randomly for experiments. Only physically intact and active individuals were selected. Adults and juveniles were separated according to their body length and secondary sexual characters. Individuals with body length \leq 3 mm and showing no development of sexual characters were defined as juveniles.

Amphipods were acclimated in 500-ml glass beakers filled with well-aerated artificial sea water (Instant Ocean®) for about 1 week before any experiments were undertaken. No more than 10 individuals were raised in each beaker. The acclimation conditions were standardized under a 14:10 h L:D regime at light density between 850 and 1200 lm m^{-2} . Temperature, salinity, pH and dissolved oxygen were maintained at $24-26^{\circ}$ C, $29-32$, $7.0-$ 8.5 and 4.7–6.8 mg 1^{-1} , respectively. These parameters were monitored every 2 days when \sim 250 ml of the cultured medium was exchanged with well-aerated artificial sea water. The amphipods were fed with four or five pieces $(1-2\text{ mm}^3)$ of freeze-dried krill after each water exchange. Two pieces of the green algal thallus *Ulva lactuca* (Linnaeus, 1753) (2 cm²) were provided as clinging substratum and supplementary food source. Any thallus piece was replaced when it was severely grazed. Faecal matter, food residue, exuviae and any undesirable algae were removed during water exchange.

Lethal tests. To investigate temperature effects, the amphipods were exposed to 0, 4, 6, 7, 8, 10, 12, 14, 20 25, 30, 31, 32, 33, 34 and 35° C at a salinity of 30 for 96 h in an environmental chamber. After acclimation, 10 amphipods were randomly selected and placed in a 500-ml beaker. A total of 50 individuals (adult or juvenile) in five replicates was tested for each temperature. Temperature was slowly adjusted from 25° C to the experimental temperature at the rate of $2-5^{\circ}C$ h⁻¹ for minimizing any drastic change of environmental conditions. No food was provided during the experimental period. All dead amphipods and exuviae were removed from the beakers during the daily inspection. The physical parameters of the medium were monitored daily during the experimental period. About 250 ml of medium was renewed every 48 h. Mortality of amphipods was recorded at the end of the experiment and the median lethal temperatures (LT_{50}) at a salinity of 30 were determined as described in the next section.

To investigate salinity effects, the amphipods were exposed to 0, 2, 5, 8, 10, 20, 30, 40, 50, 55, 58, 60, 62 and 65 salinities at 25° C for 96 h. Salinity of the medium was adjusted from 30 to the experimental value at a rate of 2–5 h^{-1} . The median lethal salinities (LS₅₀) at 25°C were also determined as described below.

Data analyses. Results from the lethal tests were analysed by probit analysis using the computer software USEPA Probit Analysis program version 1.5. Probit transformation was applied to the cumulative mortality data. The values of 96-h LT_{50} were estimated within both low (0–14 \degree C) and high (30–35 \degree C) temperature ranges, while the values of 96-h LS₅₀ were estimated within low $(0-10)$ and high (50–65) salinity ranges. Data from adults and juveniles were compared using Student's t-test (Zar 1996). The computer softwares Sigmastat version 2.03 and Sigmaplot 2000 were used.

Growth of juveniles at different temperatures and salinities

Culture. Ovigerous females were collected from the outdoor culture tanks. No more than two newly hatched juveniles were taken from each female for experiments. Ten juveniles were studied under each combination of temperature (15, 20 and 25° C) and salinity (10, 20, 30 and 40). Each juvenile was reared individually in a 100-ml beaker for any specific temperature–salinity combination. Temperature and salinity were adjusted from acclimated conditions at the rates of 2°C h^{-1} and 2 h^{-1} , respectively. Each juvenile was fed with a piece of krill $(1-2\text{ mm}^3)$ and algal thalli which had been immersed in artificial sea water of the respective combination of salinity and temperature for about an hour before the start of the experiment. The culture procedures were similar to those described in the previous section. Physical characteristics of the experimental media (temperature, salinity, pH and dissolved oxygen) were monitored every 2 days.

Growth measurement. Each juvenile was examined daily for moulting. Moulting was evidenced by the presence of the exuvium in the beaker. The day of exuvium appearance was regarded as the day of moulting. Body length (from rostrum to posterodistal end of urosomite 3 in a naturally curved posture) was measured 1 day after moulting. As the body length between the two sexes began to differ significantly at instar 7 (Tsoi 1999), growth study on juveniles was limited to the first six instars. To measure body length (BL), the

amphipod was placed on a glass slide and covered with a small sheet of thin plastic film. Each amphipod was examined for \leq 2 min to minimize the effects of hypoxia and handling stress. The image of each amphipod was recorded using a compound microscope (Nikon model SE) and a colour camera set (Teli® CCD CS 5110), and the body length was measured to 0.01 mm with a video-based image analysing program (Quantimet 500C).

Data analysis. Statistical analyses including Student's t-test, two-factor analysis of variance (ANOVA) followed by Tukey test were performed using the computer software Sigmastat version 2.03.

Results

Survival at different temperatures and salinities

At a salinity of 30, $>90\%$ of adults and juveniles survived for 96 h between 6 and 28 $^{\circ}$ C (Figure 1A). Survival was $\leq 90\%$ when the temperature was $\leq 6\degree$ C or $\geq 28\degree$ C. No amphipod survived at 0 or 35°C for >24 h. The values of the low-range 96-h LT₅₀ differed significantly between adults and juveniles (Student's t-test; $P=0.02$), indicating that adults were more tolerant to low temperature than juveniles (Table I). No amphipod remained active after 96 h of exposure to 4 and 6° C, but sluggish animals became reactivated within 15 min after transfer to 25 $^{\circ}$ C. All 'recovered' amphipods were still alive after 24 h of incubation at room temperature. There was no significant difference in the values of highrange 96-h LT₅₀ between adults and juveniles (P>0.5) (Table I).

In salinities ranging from 10 to 30, $>90\%$ of the amphipods survived for 96 h at 25°C (Figure 1B). Survival rate decreased at more extreme salinities. No amphipod could survive for $>$ 24 h and $>$ 48 h at salinities of 0 (distilled water) or 65, respectively. The values of 96-h LS_{50} between adults and juveniles were not significantly different in either low $(P=0.10)$ or high $(P=0.49)$ salinity ranges.

Growth of juveniles at different temperatures and salinities

Growth rate of juveniles expressed as increase in BL per day during the juvenile stage (instars 1–6) was calculated at different temperature–salinity combinations (Table II). Both factors affected growth rate significantly (two-factor ANOVA; $P<0.001$). Growth rate was highest at 25°C and a salinity of 20 (0.114 \pm 0.006 mm day $^{-1}$) (two-factor ANOVA with temperature and salinity as major factors; $P<0.001$). Variations in either moult increment in BL or intermoult duration, or both, could affect the growth rate of juveniles. At a salinity of 20, juveniles reared at 25° C showed the highest percentage moult increment in BL among the temperatures studied and the deviation became significant at the transition from instar 3 to 4 (Tukey test after two-factor ANOVA with temperature and instar as major factors; $P<0.05$) (Figure 2A). At 25°C, juveniles reared at a salinity of 20 showed the greatest percentage moult increment in BL among the salinities studied and the deviation became significant at the transition from instar 3 to 4 (Tukey test after two-factor ANOVA with salinity and instar as major factors; $P<0.05$) (Figure 2B).

At a salinity of 20, intermoult duration of the juveniles reared at 15° C was the longest among the temperatures studied from instar 1 (Tukey test after two-factor ANOVA with temperature and instar as major factors; $P<0.01$) (Figure 3A). In contrast, the duration

Figure 1. Survival of Hyale crassicornis adults and juveniles after 96 h of exposure to (a) different temperatures at a salinity of 30 and (b) different salinities at 25° C. Values are expressed as mean \pm SD of five replicate experiments, each of 10 individuals.

was the shortest at $25^{\circ}C$ (P<0.001). However, no significant difference was observed in intermoult duration among the four salinities studied (two-factor ANOVA with salinity and instar as major factors; $P=0.06$) (Figure 3B).

	96-h LT ₅₀ (°C)		96-h LS_{50}	
	Low temperature	High temperature	Low salinity	High salinity
Adult Juvenile	$3.2(2.6-3.6)^*$ 4.2 $(3.9-4.3)^*$	$31.7(31.1-32.3)$ $31.5(29.7-32.5)$	$2.4(1.7-2.9)$ $1.6(0.9-2.2)$	$55.5(50.9-58.1)$ $56.9(42.1 - 62.4)$

Table I. Median lethal temperature and salinity (96 h) of Hyale crassicornis juveniles and adults.

Values are expressed as mean and 95% confidence interval (in parentheses) of five replicate experiments, each of 10 individuals. *Significant difference between adult and juvenile.

Table II. Body length (BL) of *Hyale crassicornis* juveniles at instar 6, and duration and growth rate of juveniles (instars 1–6) reared at different combinations of salinity and temperature.

			Temperature $(^{\circ}C)$	
Salinity		15	20	25
10	BL (mm)	$3.40 + 0.15$	$3.16 + 0.12$	$3.09 + 0.06$
	Duration (days)	$38.5 + 3.5$	$26.5 + 3.6$	$26.9 + 3.4$
	Growth rate (mm day ⁻¹)	$0.055 + 0.005$	$0.055 + 0.045$	$0.060 + 0.003$
20	BL (mm)	$3.48 + 0.15$	$3.65 + 0.20$	$4.54 + 0.25$
	Duration (days)	$33.9 + 3.4$	$27.9 + 4.3$	$25.3 + 3.7$
	Growth rate (mm day ⁻¹)	$0.061 + 0.004$	$0.080 + 0.009$	$0.114 + 0.006$
30	BL (mm)	$2.90 + 0.16$	$3.89 + 0.18$	$3.92 + 0.18$
	Duration (days)	$33.4 + 3.5$	$30.3 + 3.6$	$29.1 + 4.8$
	Growth rate (mm day ⁻¹)	$0.044 + 0.005$	$0.081 + 0.005$	$0.009 + 0.008$
40	BL (mm)	$3.31 + 0.17$	$2.57 + 0.19$	$3.25 + 0.19$
	Duration (days)	$37.6 + 4.4$	$33.8 + 4.7$	$29.2 + 3.5$
	Growth rate (mm day ⁻¹)	$0.051 + 0.005$	$0.067 + 0.006$	$0.067 + 0.001$

Growth rate is expressed as increase of body length per day (mm day^{-1}) . Values are mean \pm SD $(n=10)$.

In summary, the growth rate of juveniles was highest at 25° C and a salinity of 20 as a result of an increase in moult increment of BL and a shortening of intermoult duration. Temperature exerted significant effects in both moult increment and intermoult duration. However, salinity only exerted its effects on moult increment, not on intermoult duration.

Discussion

Survival at different temperatures and salinities

This study shows that, on average, 90% of Hyale crassicornis could survive for 4 days at 6– 28[°]C. The species could not survive at 0 or 35[°]C. Another member of the same superfamily Talitroidea, the sub-Antarctic beach-hopper 'Orchestia' scutigerula Dana, 1853 exhibits a different range of temperature tolerance (Moore et al. 1995). The lethal temperatures of this species are $\langle -4^{\circ}$ C and \rangle 28°C. Brown and Bert (1993) suggested that the difference in temperature tolerance of crustaceans may be the consequence of adaptation to different environmental conditions. 'Orchestia' scutigerula was found beneath supralittoral stones in sub-Antarctic regions where the environmental temperature could change dramatically (Moore et al. 1995). In Hong Kong, H. *crassicornis* is exposed to water temperatures ranging between 16 and 32° C. This species exhibits a higher tolerance to high temperature but lower tolerance to low temperature than 'Orchestia' scutigerula. The difference in

Figure 2. Moult increment expressed as percentage increase in body length at each instar from 1 to 6 (mean \pm SE, $n=10$) of Hyale crassicornis juveniles reared at (a) different temperatures at a salinity of 20 and (b) different salinities at 25°C.

Figure 3. Intermoult duration (mean \pm SE, $n=10$) of each instar from 1 to 6 in *Hyale crassicornis* juveniles reared at (a) different temperatures at a salinity of 20 and (b) different salinities at 25° C.

thermal tolerance between these species is apparently associated with habitats. Thermal tolerance can also be attenuated by salinity change and the presence of certain chemicals. For instance, soft-shell snow crab Chionoecetes opilio (Fabricius, 1788) is more susceptible to thermal stress at low ambient salinity (Hardy et al. 1994). Ammonia or nitrite exposure weakens the thermal tolerance of the post-larvae of the white shrimp Penaeus setiferus (Alcaraz et al. 1997). Thus the data obtained from this study (both the high and low temperature ranges) may not reflect the survival limits of H. crassicornis in the field because of the seasonal fluctuation of salinity (heavy rainfall in summer) and pollution. Further studies are required to investigate the interacting effects of these environmental factors on the survival of this species.

Many shore invertebrates can survive in habitats where abrupt changes in salinity take place (Kinne 1971). Hyale crassicornis is able to survive ($>80\%$) in a wide range of salinities between 10 and 50. The 96-h LS_{50} values of adults at high and low salinity ranges were 55.5 and 2.4, respectively. The corresponding values for juveniles were 56.9 and 1.6. These values are similar to those reported in other euryhaline crustaceans, such as the isopod Sphaeroma serratum (Fabricius, 1787). The high and low values of 96-h LS_{50} in stage I juvenile S. serratum were 59 and 13, respectively (Charmantier and Charmantier-Daures 1994). The values of 21-day LS_{50} in the blue crab *Callinectes similis* Williams, 1966 were 60.8 and 2.6 at high and low salinity ranges, respectively (Guerin and Stickle, 1997b). The high salinity tolerance of H. crassicornis is consistent with the results from studies on various members of the same superfamily Talitroidea. For instance, the euryhaline *Traskorchestia* traskiana (Stimpson, 1857) can tolerate salinities between 2.5 and 50 for at least a week at 10° C (Koch 1991). *Orchestia gammarellus* (Pallas, 1766) survives between 1 and 60 (Moore and Francis 1986), while Transorchestia chiliensis (Milne-Edwards, 1840) survives within the range of 0.3–51 (Marsden 1980). Such tolerance is associated with their strong osmoregulatory capabilities. Spicer and Taylor (1987) reported that haemolymph sodium of O. gammarellus was actively regulated within a narrow range between 204 and 443 mmol kg⁻¹ when the amphipod was exposed to salinities of 5 (\approx 135 mmol kg⁻¹) or 40 (\approx 1160 mmol kg⁻¹).

Growth of juveniles at different temperatures and salinities

Temperature exerts a significant impact on the growth of crustaceans. The present study reveals that the growth rate of H. crassicornis is higher at 25° C than at 15° C and 20° C. Similar findings have been reported in other crustaceans. Penaeid shrimp juveniles grow faster at a higher temperature (Chen et al. 1995; Kumlu et al. 2000). The growth increment of *Penaeus vannamei* (Boone, 1931) is extremely sensitive to a moderate increase in temperature from 23 to 27°C (Wyban et al. 1995). Growth rate and moulting frequency of Penaeus japonicus Bate, 1888 are higher at 32 than 28° C (Hewitt and Duncan 2001).

Temperature also exerts effects on intermoult duration, which in turn affects the growth of crustaceans (Hartnoll 1982). Mean intermoult duration of juvenile H. crassicornis was extended from 5.08 days at 25° C to 6.79 days at 15° C. Such extension at low temperature has been frequently reported for other crustaceans. Thus, duration of larval instar in the stone crab Menippe mercenaria (Say, 1818) lengthened as temperature decreased from 30 to 25°C (Brown et al. 1992). Intermoult duration of *Penaeus merguiensis* De Man, 1888 increased from 3 days at 35° C to 12 days at 15° C (Staples and Heales 1991).

Other than temperature, salinity is another critical factor exerting significant effects on crustacean growth. For instance, dry weight of the blue crab Callinectes similis juveniles

reared at a salinity of 5 was half of those reared at 10 (Guerin and Stickle 1997b). In this study, we found that the optimal salinity for growth of H. crassicornis juveniles is 20 at 25 $^{\circ}$ C. The phenomenon can be explained by the physiological relationship between salinity and metabolism. A higher oxygen consumption rate may indicate a higher energy requirement for osmoregulation under osmotic stress, reducing energy available for growth (Einarson 1993). The fasting and feeding oxygen consumption of P. vannamei juveniles increases under low ambient salinity because more energy is diverted to maintain homeostasis (Rosas et al. 2001). The scope for growth in the blue crab Callinectes similis juveniles is the highest at a salinity of 35 (Guerin and Stickle 1997b). Penaeus chinensis exhibits the fastest growth rate at the isosmotic point of 16.5, suggesting that less energy is spent on osmoregulation at this salinity (Chen et al. 1992, 1996). Other than energy consumption, utilization of nutrients is also affected by salinity, leading to altered growth rate (Venkataramiah et al. 1972).

Any effects leading to either an increase in moult increment of body size or shortening in intermoult duration could result in a higher growth rate. Our study reveals that H. crassicornis juveniles reared at a salinity of 20 had the highest percentage moult increment in body length compared with those reared at 15 and 25. Yet salinity exerts little effect on intermoult duration of the juveniles. This phenomenon has also been reported in the banana shrimp P. merguiensis (Staples and Heales 1991). Salinity affects growth of this shrimp through changes in the moult increment rather than intermoult duration. Insignificant effects of salinity on intermoult duration have been reported in juvenile Callinectes sapidus Rathbun, 1896 (Cadman and Weinstein 1988; Guerin and Stickle 1992), Menippe mercenaria (Brown et al. 1992) and P. chinensis (Chen et al. 1992). Yet some crustaceans exhibit prolonged developmental duration only under extreme salinity. Examples include Rhithropanopeus harrisii (Gould, 1841) post-larvae at salinities of 2.5 and 40 (Costlow et al. 1966) and Armases miersii (Rathbun, 1897) larvae and juveniles at salinities of 5 and 45–55 (Anger 1996). This is consistent with our observation that salinity has little influence on intermoult duration.

Environmental factors interact to produce physiological effects on marine invertebrates (Staples and Heales 1991; O'Brien 1994). These effects in turn affect growth rate and growth pattern of crustaceans (Hartnoll 2001). The highest growth rate of H. crassicornis juveniles was 0.11 mm day⁻¹ at 25°C and a salinity of 20, while the growth rate of animals reared at other temperature–salinity combinations ranged between 0.04 and 0.08 mm day⁻¹. Such growth rates are comparable to those reported in other amphipods. For instance, the growth rates of Hyale barbicornis Hiwatari and Kajihara, 1981 juveniles at 24° C were 0.10 and 0.15 mm day⁻¹ in males and females, respectively (Hiwatari and Kajihara 1988). The optimal combination of conditions for growth of H. crassicornis juveniles was 25° C and a salinity of 20. The optimal temperature–salinity combination for growth is species-specific (Chen and Chen 1993; Kumlu et al. 2001). For example, the optimal culture conditions in juvenile or larval penaeid shrimps: P. semisulcatus De Haan, 1844, P. esculentus Haswell, 1879, P. merguiensis and P. chinensis are 30°C and 30–35 salinity (Kumlu et al. 2000), 30°C and 30 salinity (O'Brien 1994), 28[°]C and 28 (Staples and Heales 1991) and 30[°]C and 20–30 salinity (Chen et al. 1996), respectively.

In summary, the growth pattern of H. *crassicornis* is controlled by both temperature and salinity. Change in moult increment of body size is driven by both temperature and salinity while intermoult duration appears to be influenced by temperature alone. Growth of H. crassicornis juveniles is optimal at 25° C and a salinity of 20. While these data from our laboratory experiments are valuable in the development of pollution bioassays, further

studies are required to understand the impact of temperature and salinity variations on natural populations of this amphipod.

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