Enhanced growth of juvenile *Tachypleus tridentatus* (Chelicerata: Xiphosura) in the laboratory: a step towards population restocking for conservation of the species

Yan Chen1,2, C. W. Lau1, S. G. Cheung1,3, C. H. Ke4, Paul K. S. Shin1,3,*

1Department of Biology and Chemistry, and 3State Key Laboratory in Marine Pollution, City University of Hong Kong, Tat Chee Avenue, Kowloon, Hong Kong SAR

2College of Marine Sciences, Shanghai Ocean University, No. 999, Hu Cheng Loop-road, Lingang New City, Shanghai, PR China

4State Key Laboratory of Marine Environmental Science, College of Oceanography and Environmental Science, Xiamen University, Xiamen, Fujian, PR China

ABSTRACT: Juveniles of artificially bred *Tachypleus tridentatus* were reared in the laboratory for 15.5 mo. Fast growth (at least 3 times faster than in previous studies) and low mortality were observed, with juveniles reaching the 9th instar stage and having a cumulative mortality of 61.3% at the end of the rearing period. Such enhanced growth can potentially be attributed to high seawater temperature, sufficient living space and constant water flow. Isometric growth of prosomal width and other growth parameters including opisthosoma width, eye distance and prosomal length was found. Two distinct growth phases were detected for the opisthosoma length, with zero growth from the 1st to 2nd instar but isometric growth from the 2nd to 9th instar stage. Growth in telson length was positively allometric from the 1st to 3rd instar and isometric from the 3rd to 9th instar stage. The increase in wet weight and prosomal width was isometric throughout the 9 developmental stages. Based on the allometric growth patterns, the juveniles can reach adult size within 4 yr. The results may serve as a basis for the large-scale culture of this species for the purposes of restocking and restoration of natural populations.

KEY WORDS: Horseshoe crab · *Tachypleus tridentatus* · Juvenile growth · Allometry

RESALE OR REPLICATION NOT PERMITTED WITHOUT WRITTEN CONSENT OF THE PUBLISHER

INTRODUCTION

Horseshoe crabs are an archaic group of marine chelicerates, with the oldest fossils dating back to the Upper Ordovician 445 million years ago (Rudkin et al. 2008). There are 4 living horseshoe crab species, including one American species, *Limulus polyphemus* (Linnaeus, 1758), and 3 Asian species, *Carcinoscorpius rotundicauda* (Latreille, 1802), *Tachypleus tridentatus* (Leach, 1819) and *T. gigas* (Müller, 1785). Horseshoe crab populations are rapidly declining around the world, as has been reported in the USA (Botton & Haskin 1984, Bell & Henderson 1993, Shuster et al. 2003, Smith et al. 2009), Japan (Itow 1998), Taiwan (Chen et al. 2004, Hsieh & Chen 2009), Singapore (Hong 2004) and the Philippines (Schoppe 2002). Horseshoe crabs were known to occur in great numbers in Hong Kong some 30 yr ago, but now, they can only be found in much lower numbers (Chiu & Morton 1999, 2003, Li 2008). An updated survey, using both random transect and walk-through search methods at 17 shores in Hong Kong in summer and winter showed that the density of juvenile horseshoe crabs *T. tridentatus* and *C. rotundicauda* had declined significantly (by over 90%) since 2002 (Shin et al. 2009). The loss of breeding habitats, coupled with the slow growth of these crabs, means that these living fossils may be facing a potential threat of extinction.
Use of captive breeding for population enhancement in animal conservation has increased markedly in the last decade (Olney et al. 1994). In Taiwan, more than 10,000 individuals of second-instar juvenile *Tachypleus tridentatus* produced artificially in the laboratory were released in the protected area in Kinmen in 2002 (Chen et al. 2004), while about 40,000 juveniles were released in the wild in Xiamen, China, in the same year. A successful restocking program relies heavily on an adequate supply of juveniles from artificial breeding, as juveniles released in the wild may suffer high mortality from predation, diseases, food availability, etc. The problems with juvenile horseshoe crabs in culture include low fertilization success, high juvenile mortality and slow growth of the instars (Sekiguchi et al. 1988, Chatterji 1994). The most extensive study of horseshoe crab growth was conducted by Sekiguchi (1988) on the juveniles of *Limulus polyphemus* and *T. tridentatus*, which required 9 yr to develop into 14th and 10th instars, respectively. The study itself, however, may not be representative because of both low numbers of specimens and survivors in the study, as an initial number of 44 and 33 ind. were used for *L. polyphemus* and *T. tridentatus*, respectively, and only one individual of each species survived at the end of the experiment.

The actual life span and exact number of eclosures events in horseshoe crabs are unknown (Tanacredi 2001, Carmichael et al. 2003, Lee & Morton 2005), as it is difficult to accurately assess age in horseshoe crabs (Botton & Ropes 1988). Goto & Hattori (1929) suggested that *Tachypleus tridentatus* molts 12 to 13 times before maturity, while Kawahara (1984) proposed that males molt 14 times in 9 yr and females 15 times in 10 yr. Sekiguchi et al. (1988), on the other hand, reported that *T. tridentatus* males molted 15 times within 14 yr, based on an allometric growth pattern identified in laboratory culture over a period of 9 yr. Asano (1942), however, reported that the species reached maturity in 15 to 16 yr after 17 or 18 eclosures based on field observations and measurements of exuvial sizes. To follow individuals’ growth from hatching to maturity in a natural habitat is challenging because it is impractical to produce a long-lasting external tag for juvenile horseshoe crab due to their regular eclosures (Sekiguchi et al. 1988, Lee & Morton 2005). Therefore, indirect methods including examination of either living individuals (Carmichael et al. 2003) or exuvias (Shuster 1958, 1982) have been used. Another way to obtain growth data is by studying individuals in culture (Sekiguchi 1988). By following the growth and size of individual larval instars and assuming that individual growth between larval instars is the same in culture as in the wild, results may be extrapolated to estimate size and age of individuals in the field.

The main aim of the present study was to investigate growth of the Chinese horseshoe crab *Tachypleus tridentatus* in culture and assess the possibility of producing culture conditions which will permit successful large-scale production of this species for population enhancement in the wild.

**MATERIALS AND METHODS**

**Artificial insemination.** A mating pair of horseshoe crabs *Tachypleus tridentatus* was purchased from a fish market, and artificial insemination and breeding was carried out on 23 August 2007. The ventral sides of the prosoma of the mating pairs were cut open. Eggs were collected using a spoon, while sperm was collected using a syringe. Eggs were washed with seawater and *in vitro* fertilization was carried out by mixing the eggs with diluted sperm (diluted 4 times using filtered natural seawater). The fertilization rate was over 95\%, and the fertilized eggs were then incubated in the laboratory.

**Crab maintenance.** A fiber glass tank (200 × 100 × 25 cm) equipped with a water re-circulating system was used for the culturing of fertilized eggs and subsequent juveniles. The re-circulating system consisted of a protein skimmer, ceramic rings, UV light and filter cottons. Natural seawater (30‰) collected from a clean site along the east coast of Hong Kong was used in the culture process and maintained at 28 to 30°C by electric heaters. The water flow was maintained at 12 l min\(^{-1}\). Half of the water in the tank was renewed every week, and air stones were put inside the tank to provide aeration. The dark:light cycle was set at 12 h dark:12 h light.

Prior to hatching, the fertilized eggs were suspended in running water inside the tank using small baskets to maintain a high level of oxygen around the eggs, whereas after hatching, the juveniles were cultured directly in the tank with running water and coral sand (0.5 to 2.0 mm in particle size, 5 cm in depth), as juvenile horseshoe crabs prefer burrowing into the substrate.

No food was provided to the trilobite larvae as they do not feed (Carmichael et al. 2009). Starting from the 2nd larval stage, juveniles were fed daily at night. The 2nd and 3rd instars were fed with newly hatched larval brine shrimp *Artemia salina* which was left in the tank overnight, and unconsumed shrimp were removed the following morning. During feeding, water circulation was stopped but was resumed the next morning. The 4th and 5th instars were fed with defrosted adult brine shrimps *A. salina*, whereas the 6th instars and beyond were fed with the meat of clam *Ruditapes philippinarum* and shrimp *Metapenaeus ensis*, which were chopped into small pieces. The food was left in the
tank overnight, and unconsumed food was removed the following morning. Water circulation was maintained when the juveniles were fed.

**Growth measurements and data analysis.** The morphometric parameters measured are shown in Fig. 1. Parameters were measured weekly at earlier stages and monthly at later stages. The condition of the horseshoe crabs was checked daily. Deceased individuals as well as exuviae were collected and the number recorded every day. To avoid handling stress, especially for small-sized individuals, exuviae collected during the rearing period were used for the detailed size measurements of various body parts in order to construct growth models. Fifty specimens were measured from the 1st to the 7th instar, whilst 28 specimens were used for the 8th and 9th instars. Measurements of small-sized individuals, i.e. the 1st, 2nd and 3rd instars, were conducted under a stereomicroscope.

The allometric growth of each parameter \( y \) was expressed as a power function of prosomal width \( x \) (Fuiman 1983) in the form of an equation \( y = ax^b \). A \( b \) coefficient was estimated from the linear regression of an allometric growth curve, where \( \log y = \log a + b \log x \). This coefficient identifies the allometric growth pattern of *Tachypleus tridentatus* such that values of \( b = 1 \) for length and \( b = 3 \) for weight indicate isometric allometry; values of \( b > 1 \) for length and \( b > 3 \) for weight indicate positive allometry, and values of \( b < 1 \) for length and \( b < 3 \) for weight indicate negative allometry (Sekiguchi et al. 1988, Osse & van den Boogaart 2004, Lee & Morton 2005). Inflexion points of the growth curves were determined using the iteration procedure according to van Snik et al. (1997).

**RESULTS**

**Mortality**

A low hatching rate (0.79%, or 155 ind. from some 20,000 eggs) was recorded. However, survival rates were high at subsequent instar stages. The highest mortality (32.7%) was observed for the 1st instar, whereas mortality decreased to <20% for both the 2nd and 3rd instars, <5% for the 4th to 9th instars and 0% for the 6th, 8th and 9th instars (Table 1). The cumulative mortality from the 1st to 9th instar was 61.3%.

**Developmental stages**

In total, 155 trilobite larvae were hatched from 2 October to 5 November 2007, and the hatching time varied between 39 and 74 d after fertilization, or 55.2 d on average. The average age recorded for each developmental stage is presented in Table 1. The lowest growth was recorded for a single individual which remained as a 1st instar for almost 11 mo before developing into a 2nd instar, when most of the individuals had already reached the 7th instar stage. This individual developed into a 3rd instar at the age of 13.5 mo, and a 4th instar at the age of 16 mo, when most of the individuals were already 9th instars.

**Growth**

**Allometric growth of size and weight**

Mean values (±SD) of each morphometric parameter for each instar stage are summarized in Table 1, and the allometric relationships are shown in Fig. 2. The growth of the eye distance \( a = 0.6201, b = 0.9591, \)**
Table 1. *Tachypleus tridentatus*. Growth (mean ± SD) of juvenile horseshoe crabs from the 1st to 9th instar stage. n = 50 for each of the size/weight measurement of individuals from the 1st to 7th instar stage; n = 28 for the measurements of individuals from the 8th and 9th instar stage.

<table>
<thead>
<tr>
<th>Instar stage</th>
<th>Age (d)</th>
<th>Mortality (%)</th>
<th>Proosomal width</th>
<th>Eye distance</th>
<th>Opisthosoma width</th>
<th>Proosomal length</th>
<th>Opisthosoma length</th>
<th>Telson length</th>
<th>Wet weight (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>53.2 ± 4.3</td>
<td>32.7</td>
<td>6.26 ± 0.16</td>
<td>3.82 ± 0.12</td>
<td>4.79 ± 0.09</td>
<td>3.78 ± 0.16</td>
<td>2.88 ± 0.13</td>
<td>0</td>
<td>0.02 ± 0.002</td>
</tr>
<tr>
<td>II</td>
<td>133.9 ± 7.7</td>
<td>18.4</td>
<td>8.86 ± 0.32</td>
<td>4.89 ± 0.18</td>
<td>6.39 ± 0.21</td>
<td>5.30 ± 0.18</td>
<td>2.88 ± 0.16</td>
<td>3.55 ± 0.32</td>
<td>0.06 ± 0.01</td>
</tr>
<tr>
<td>III</td>
<td>154.6 ± 8.6</td>
<td>15.6</td>
<td>12.3 ± 0.53</td>
<td>6.64 ± 0.19</td>
<td>8.67 ± 0.33</td>
<td>7.41 ± 0.25</td>
<td>4.02 ± 0.13</td>
<td>8.05 ± 0.45</td>
<td>0.14 ± 0.02</td>
</tr>
<tr>
<td>IV</td>
<td>198.7 ± 13.1</td>
<td>3.3</td>
<td>16.4 ± 0.69</td>
<td>8.83 ± 0.29</td>
<td>11.7 ± 0.44</td>
<td>9.90 ± 0.39</td>
<td>5.41 ± 0.20</td>
<td>13.0 ± 0.82</td>
<td>0.32 ± 0.03</td>
</tr>
<tr>
<td>V</td>
<td>233.3 ± 8.2</td>
<td>1.7</td>
<td>22.0 ± 1.25</td>
<td>12.0 ± 0.66</td>
<td>16.0 ± 0.98</td>
<td>13.5 ± 0.80</td>
<td>7.32 ± 0.45</td>
<td>18.8 ± 2.01</td>
<td>0.72 ± 0.06</td>
</tr>
<tr>
<td>VI</td>
<td>288.8 ± 9.9</td>
<td>0</td>
<td>28.9 ± 1.52</td>
<td>20.7 ± 1.14</td>
<td>28.0 ± 1.75</td>
<td>24.0 ± 1.35</td>
<td>13.3 ± 0.73</td>
<td>32.0 ± 3.53</td>
<td>4.43 ± 0.73</td>
</tr>
<tr>
<td>VII</td>
<td>335.1 ± 9.2</td>
<td>1.9</td>
<td>38.3 ± 2.49</td>
<td>20.7 ± 1.14</td>
<td>35.9 ± 2.52</td>
<td>31.4 ± 2.22</td>
<td>17.6 ± 1.04</td>
<td>44.5 ± 4.67</td>
<td>9.33 ± 1.56</td>
</tr>
<tr>
<td>VIII</td>
<td>385.4 ± 12.6</td>
<td>0</td>
<td>49.8 ± 3.35</td>
<td>26.7 ± 1.60</td>
<td>47.0 ± 3.77</td>
<td>44.6 ± 3.71</td>
<td>23.3 ± 1.92</td>
<td>61.4 ± 6.09</td>
<td>20.8 ± 4.17</td>
</tr>
<tr>
<td>IX</td>
<td>463.5 ± 7.5</td>
<td>0</td>
<td>65.7 ± 6.09</td>
<td>35.2 ± 2.77</td>
<td>61.4 ± 6.09</td>
<td>47.0 ± 4.03</td>
<td>23.3 ± 1.92</td>
<td>61.4 ± 6.09</td>
<td>20.8 ± 4.17</td>
</tr>
</tbody>
</table>

Growth estimations

Horseshoe crabs show stepwise growth (Shuster 1954, Shuster & Sekiguchi 2003); therefore, the size increment during their intermolt periods is barely discernible, and the prosomal width is always correlated with the development stage as well as with age (Lee & Morton 2005). Regressions relating prosomal width and body wet weight and instar stage are shown in Fig. 3a and that relating age and instar stage is shown in Fig. 3b. The corresponding equations are:

\[
L = 0.4982 e^{0.2905t} \quad (r^2 = 0.9986)
\]
\[
Wt = 0.0106 e^{0.8429i} \quad (r^2 = 0.9986)
\]
\[
T = 3.145 e^{0.1791i} \quad (r^2 = 0.9955)
\]

where \(L\) represents prosomal width in cm, \(Wt\) represents wet weight in g, \(T\) represents age in month, and \(i\) represents the instar stage.

Growth data collected from the 1st to 9th instars were used to estimate the stepwise growth of horseshoe crabs beyond the 9th instar stage (Fig. 3a). The estimated ages of instars beyond Stage 9, however, were based on the results obtained from the 2nd to 9th instars, as the age–instar stage relationship for the 1st instar was different from other stages (Fig. 3b). Growth increased about one-third with each molt. According to the extrapolated data, these horseshoe crabs could develop into 11th instars in 2 yr, with a prosomal width of 12.2 cm and wet weight of 113 g. Two more ecdyses would occur in both the third and fourth year, with a prosomal width of 38.9 cm and wet weight of 3.3 kg for a 15th instar.

Developmental deformities

All individuals from the 1st to 4th instar stage possessed perfectly symmetrical prosoma. Three individuals at the 5th instar stage (4.3%) were found to have either a spiral or shorter telson. The number of individuals with such deformities increased to 11 (16%) at the 6th instar stage. A partial concave margin of the prosoma was recorded in 5 ind. after the 6th instar stage. Degradation of a single eye was observed in 3 ind. (2 on the right eye) after the 7th instar stage (4.3%). All these individuals behaved normally, but these deformities remained during the study period.

DISCUSSION

Mortality

In the present study, a very low hatching success was recorded. This could possibly be due to fungal infection at the incubation stage of the fertilized eggs in the holding baskets within the re-circulating water tank (Schreibman & Zarnoch 2009). However, more than one-third of the 1st instar horseshoe crabs survived to 9th instar stage (cumulative mortality 61.3%). The survival rate was higher than that of a wild population of the American species *Limulus polyphemus* in Pleasant
Fig. 2. *Tachypleus tridentatus*. Allometric growth equations and relationship between the prosomal width with different body part sizes, and with wet weight of horseshoe crab juveniles from the 1st to 9th instar stage. (a) Eye distance; (b) opisthosa width; (c) prosomal length; (d) opisthosa length; (e) telson length (no telson at the 1st instar stage); (f) wet weight. (n = 50 for each measurement of individuals from the 1st to 7th instar stage; n = 28 for individuals from the 8th and 9th instar stage)
Bay, USA, where the cumulative mortality was 99.6% from the 1st to the 7th instar stage (which lasted 1 yr) (Carmichael et al. 2003). The mortality at each stage in the Carmichael et al. study was also higher in *L. polyphemus* (58, 80, 27, 63, 67, and 36% at the 1st, 2nd, 3rd, 4th, 5th and 7th instar stage, respectively) than in the *Tachypleus tridentatus* examined in our study. A lower mortality was also obtained for laboratory-hatched and -cultured *Carcinoscorpius rotundicauda* and *T. gigas* for which the mortality after 1 yr was 39.9 and 34.5%, respectively, with the abundant instar stages being the 6th instar and 5th instar, respectively (Zadeh et al. 2009). According to Carmichael et al. (2003), mortality beyond the 8th instar stage was rare, which is consistent with our study in which zero mortality was obtained for both the 8th and 9th instars. The elevated survival rate under laboratory conditions at the early developmental stages observed in the present study can be significant for the conservation of this species, as a large number of juveniles can be obtained from a small number of adult horseshoe crabs for a restocking program, especially important in areas where adult horseshoe crabs are endangered.

**Developmental stages**

In the present study, large variations in hatching times were noted. Similar observations were reported by Zadeh et al. (2009), in which some laboratory-cultured *Tachypleus gigas* and *Carcinoscorpius rotundicauda* larvae survived the 1st instar stage but did not molt for 170 d. Such delayed molting is also evident in wild *Limulus polyphemus* populations where larvae emerged from the sandy nests without having molted during winter (Botton et al. 1992). In the present breeding experiment all full siblings were cultured together. Our results were obtained from only one pair of adult *T. tridentatus*, and presumably there was little generic variation among them. Mishra (2009) postulated that the variability in hatching time might be associated with a molting protein in horseshoe crabs. A lower protein level would delay subsequent molting events, leading to slow-developing outliers. It is likely that the molting protein in question is the molting hormone 20-hydroxyecdysone (Shuster & Sekiguchi 2003). However, further studies are required to elucidate the involvement of such a protein and other physiological events related to molting changes in horseshoe crab development.

**Growth**

There are very few records of consecutive data on the stepwise growth of horseshoe crabs, due to the difficulties in raising them from eggs to adults in the laboratory (Sekiguchi et al. 1988, Chatterji 1994, Shuster & Sekiguchi 2003). A comparison of the prosomal width of different instars is shown in Table 2. Sizes of instars obtained from field and laboratory studies by Chen et al. (2010) were similar to ours, but those obtained in a laboratory study by Sekiguchi (1988) were smaller.

Unusually fast growth was recorded in the present study. In contrast to general growth events with 6 ecdyses in the first 3 yr after hatching (Rudloe 1981,
Shuster 1982, Mikkelsen 1988, Sekiguchi et al. 1988, Carmichael et al. 2003) and 1 ecdysis once a year thereafter until maturation (Shuster 1954, 1958, 1982, Mikkelsen 1988, Sekiguchi 1988, Chatterji 1994), juvenile *Tachypleus tridentatus* in our study experienced 6 ecdyses within a year (including approximately 2 mo before hatching), and 2 more ecdyses in the following 4 mo. Such fast growth can be attributed to laboratory rearing with high water temperatures and an adequate food supply.

Chiu & Morton (2004) reported the seasonal pattern of feeding behaviour of juvenile horseshoe crabs at an intertidal zone in Hong Kong. The highest number of feeding trails was recorded in summer, while only a few juveniles were seen in winter, when the temperature can be <10°C, implying that horseshoe crabs barely eat at low temperatures in the field. In contrast, laboratory-cultured juveniles in our study were maintained at 28 to 30°C and fed every day. A slower growth in horseshoe crab juveniles at lower temperatures has also been reported in previous studies (Shuster 1982, Yeh 1999), which showed that ecdysis continued when the temperature remained at >28°C but halted at <22°C. Low temperatures reduce molting hormone levels to a critical level in *Limulus polyphemus* and hence suspend the ecdysis at temperatures <20°C (Jegla & Costlow 1982).

Chen et al. (2004) suggested that 28 to 31°C is the most suitable temperature for incubating eggs and rearing juveniles. The present study showed that the growth of *Tachypleus tridentatus* could proceed throughout the year, provided that the seawater temperature was optimal (28 to 30°C). A much slower growth for *T. tridentatus*, however, was obtained by Sekiguchi et al. (1988). In the first year of rearing, the juvenile *T. tridentatus* remained as 1st instars when reared at room temperature but developed into 2nd instars when reared at 30°C. Altogether 5 ecdyses (i.e. the 6th instar stage) were obtained in the first 3 yr; thereafter, there was 1 ecdysis once a year until the crabs reached the 10th instar stage in the seventh year. Such differences might be attributed to differences in rearing conditions, such as living space and water quality. Juveniles in the present study were maintained in a container (200 × 100 × 25 cm) much greater in size than those used in the study of Sekiguchi et al. (1988), in which the size of containers varied from 3.4 × 3.4 × 3 cm to 36 × 25.5 × 10 cm, depending on the size of the juveniles. In addition, aerated seawater was used in the present study and the water was re-circulated. In contrast, no circulation or aeration was provided in Sekiguchi et al.’s (1988) study and seawater was changed daily.

Nevertheless, under the same rearing conditions as were used for *T. tridentatus*, a much faster growth was recorded for juvenile *Limulus polyphemus* (Sekiguchi et al. 1988) with 5, 3, 2 and 1 ecdyses in the first, second, third and fourth year, respectively.

Based on the data collected on the stepwise growth of the first 9 instar stages of *Tachypleus tridentatus*, the present study revealed isometrical growth (*b* ≈ 1) between prosomal width and other growth parameters including eye distance, opisthosaoma width, prosomal length and opisthosaoma length (except for the zero growth of opisthosaoma length between the 1st and 2nd instar). Rapid elongation of the telson was observed at the initial growth phase (*b* > 1 before the 4th instar stage) but became isometric at later growth phases (*b* ≈ 1 after the 4th instar stage). Therefore, with the exception of the telson, the shape of horseshoe crab at different developmental stages is similar. This is consistent with the results obtained by Sekiguchi et al. (1988) on *T. tridentatus* and *Limulus polyphemus*. An increase in the wet weight with the prosomal width was isometric (*b* = 3) in the present study. A similar growth pattern of wet weight (*b* = 2.9682) was obtained by Lee & Morton (2005) based on 6 consecutive instars of juvenile *T. tridentatus* which varied between 20 and 100 mm in prosomal width.

Based on the allometric growth patterns identified in the present study, *Tachypleus tridentatus* could develop into the 14th and 15th instar in 4 yr, with a corresponding prosomal width of 29.1 and 38.9 cm and a wet weight of 1.4 and 3.3 kg, respectively. Such predicted growth information is consistent with data obtained by Chiu & Morton (2003) from a wild *T. tridentatus* population in Hong Kong.

**Developmental deformities**

Deformities observed in juvenile horseshoe crabs in the present study may not be attributed to any genetic mutation or genetic diseases, since all the symptoms

<table>
<thead>
<tr>
<th>Year</th>
<th>Instar stage</th>
<th>Age (mo)</th>
<th>Prosome width (cm)</th>
<th>Wet weight (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>I—VII a</td>
<td>11.2</td>
<td>3.8</td>
<td>3.8</td>
</tr>
<tr>
<td>2</td>
<td>VIII—IX a</td>
<td>15.5</td>
<td>6.6</td>
<td>19.3</td>
</tr>
<tr>
<td></td>
<td>X b</td>
<td>18.9</td>
<td>9.1</td>
<td>48.5</td>
</tr>
<tr>
<td></td>
<td>XI b</td>
<td>22.6</td>
<td>12.2</td>
<td>113</td>
</tr>
<tr>
<td>3</td>
<td>XII b</td>
<td>27.0</td>
<td>16.3</td>
<td>262</td>
</tr>
<tr>
<td></td>
<td>XIII b</td>
<td>32.3</td>
<td>21.8</td>
<td>608</td>
</tr>
<tr>
<td>4</td>
<td>XIV b</td>
<td>38.6</td>
<td>29.1</td>
<td>1413</td>
</tr>
<tr>
<td></td>
<td>XV b</td>
<td>46.2</td>
<td>38.9</td>
<td>3283</td>
</tr>
</tbody>
</table>

*a*Data recorded from the present study  
*b*Data estimated from stepwise growth equations obtained in Fig. 3

Table 2. *Tachypleus tridentatus*. Estimation of the stepwise growth of the horseshoe crabs
were observed only in elder juveniles beyond the 5th instar stage. Suboptimal rearing conditions including a very restricted diet and small rearing space may be possible contributing factors. Juvenile horseshoe crabs in the wild are benthic feeders and subsist mainly on polychaetes, oligochaetes, crustaceans, bivalves and gastropods (Carmichael et al. 2004, Zhou & Morton 2004). The restricted diet provided for the laboratory-cultured juveniles included only shrimp and clam meat, which might not provide a balanced diet for the rapid growth of the juveniles. Using carbon and nitrogen stable isotopes as tracers, juveniles were found to use a diet of mixed composition, which changed with horseshoe crab size, and contained potentially high concentrations of particulate organic matter (Carmichael et al. 2009). In some cases, a spiral or shorter telson was found and possibly attributed to the rectangular shape of the culturing tank. Some individuals stayed in the corner of the tank when they tried to shed their exuvias during ec dysis. Their telsons, therefore, could not be straightened but became spiral or even broken. It has been suggested that limited space, handling practices or insufficient water flow can encumber molting or increase exposure to and infection by bacteria or other parasites (Smith & Berkson 2005). Further investigations are needed to determine the optimal diet and space requirements for growth.

**Culture for captive release**

Horseshoe crabs are valued for ecological, economic and educational purposes. The rapid decline in horseshoe crab populations around the world and especially in Asia, however, is evident and becoming a growing concern. A successful restocking program to enhance natural populations relies heavily on an adequate supply of juveniles for release. Several captive release or induced spawning programs have been reported in China (Hong et al. 2009), Japan (see overview by Tsuchiya 2009) and Taiwan (Chen et al. 2009) for *Tachypleus tridentatus* and in the USA for *Limulus polyphemus* (e.g. Cuomo 2009, Schreibman & Zarnoch 2009, Tzafrir-Prag et al. 2009). Of these programs, the captive release effort at the Kasaoka Municipal Horseshoe Crab Museum, Japan, has a long history (Tsuchiya 2009). Juveniles *T. tridentatus* were reared in tanks with mud and water, and the water was exchanged either once every 20 d or once a month. The crabs were fed brine shrimp, TetraMin, chopped clams and worms over their growth period. It was reported that a hatched juvenile developed into an adult in its 11th year (Tsuchiya 2009). Hong et al. (2009) also showed that juveniles grow and survive better in the presence of sand as compared to mud in the seawater tanks, and Cuomo (2009) and Tzafrir-Prag et al. (2009) demonstrated that food quality, type and digestibility have a strong effect on post-hatch molts in laboratory culture. The present study has demonstrated a much enhanced growth of *T. tridentatus* in laboratory culture, possibly due to high seawater temperatures, a large culture space and constant water flow. The results may serve as a basis for the large-scale culture of the species for restocking and restoration of natural populations.

**Acknowledgements.** This study was funded by the Ocean Park Conservation Foundation of Hong Kong.

**LITERATURE CITED**


Hong RF (2004) Population and distribution of horseshoe crab *Carcinoscorpius rotundicauda* at the Kranji Nature rail estuaries, Western Johor Straits. Project Report 49. The National University of Singapore, Department of Biological Sciences, Singapore


Li HY (2008) The conservation of horseshoe crabs in Hong Kong. MPhil thesis, City University of Hong Kong, Hong Kong


Shuster CN Jr (1958) ‘Study these’, the story of the horseshoe ‘crab’. Wilmington Public Schools Staff Reporter 10:4–5


van Snik GMJ, van den Boogaart JGM, Osse JWM (1997)
Larval growth patterns in *Cyprinus carpio* and *Clarias gariepinus* with attention to fin fold. J Fish Biol 50: 1339–1352

Yeh HY (1999) Life cycle, juvenile habitat and conservation strategies of the horseshoe crab (*Tachypleus tridentatus*) in Kinmen. MPhil thesis, National Taiwan University, Taiwan (in Chinese)


Editorial responsibility: Hans Heinrich Janssen, Oldendorf/Luhe, Germany

Submitted: April 8, 2010; Accepted: September 6, 2010
Proofs received from author(s): October 27, 2010