Effects of the brown algae *Sargassum horneri* and *Saccharina japonica* on survival, growth and resistance of small sea urchins *Strongylocentrotus intermedius*

Fangyuan Hu, Mingfang Yang, Peng Ding, Xu Zhang, Zhouling Chen, Jingyun Ding, Xiaomei Chi, Jia Luo, Chong Zhao*, & Yaqing Chang*

Mass mortality of the long line culture of the sea urchin *Strongylocentrotus intermedius* in summer, which is greatly associated with their disease, energy storage and resistant abilities, is the most serious problem for the development of the aquaculture. Here, a feeding experiment was conducted for ~9 weeks to investigate the survival, growth and gonadal development of small *S. intermedius* (~3 cm) fed either brown algae *Sargassum horneri* or *Saccharina japonica*. Subsequently, we assessed their resistant abilities via observing the behaviors of righting, tube feet extension and Aristotle’s lantern reflex at both moderately elevated and acutely changed water temperatures. Sea urchins fed *S. horneri* showed significantly fewer diseased individuals and slower gonadal development than those fed *S. japonica*. Consistently, significantly greater Aristotle’s lantern reflex occurred in sea urchins fed *S. horneri* at moderately elevated temperatures. These findings suggest that *S. horneri* has direct application potential as food for the long line culture of *S. intermedius* in summer because of the advantage in health, energy storage (avoid the energy loss caused by gonadal development at small body sizes) and resistance abilities. In comparison, sea urchins fed *S. japonica* outperformed those fed *S. horneri* for all experimental behaviors under the acutely changed water temperatures. These findings clearly suggest that *S. intermedius* fed *S. japonica* is more suitable for the areas with cold water mass in summer, because it can effectively avoid or reduce the negative impacts of acute changes of water temperature on sea urchins. The present study provides valuable information into the management of the long line culture of *S. intermedius* in summer.

The sea urchin *Strongylocentrotus intermedius* is a seafood delicacy with high commercial value in the world. Interest in *S. intermedius* aquaculture has grown rapidly in the last decade. The annual production of sea urchins, for example, was over 8,844 tons in China in 2018. Long line culture is the most important approach to meeting market demands for *S. intermedius*. The brown alga *Saccharina japonica* is commonly used as food for *S. intermedius* aquaculture in Japan and northern China. However, fresh *S. japonica* is largely unavailable in summer and thus dried *S. japonica* is used for food at that season. Our previous study found that the brown alga *Sargassum horneri* is an alternative diet for *S. japonica* and consequently addressed the food shortage problem for the aquaculture of *S. intermedius* in summer. However, mass mortality of the long line culture of *S. intermedius* in summer, which is greatly associated with their disease, energy storage, resistant abilities, is the most serious unaddressed problem for the aquaculture of *S. intermedius*.

Reproduction greatly consumes stored energy and is probably associated with various diseases even death in commercially important marine invertebrates such as the oyster *Crassostrea gigas*. Cultured *S. intermedius* (~3 cm of test diameter) showed precocious gonads in July in northern China, suggesting that the excessive allocation of energy to gametogenesis may be responsible for the mortality of sea urchins in summer.

Key Laboratory of Mariculture & Stock Enhancement in North China’s Sea, Ministry of Agriculture, Dalian Ocean University, Dalian 116023, China. *Email: chongzhao@dlou.edu.cn; changlab@hotmail.com
Strongylocentrotus intermedius fed S. horneri showed significantly lower gonad yield than those fed S. japonica in summer. Low gonad production commonly results in less gametogenesis in sea urchin gonads. Therefore, S. horneri is probably valuable to prevent precocious puberty of S. intermedius and subsequently to enhance the energy storage.

In addition, long line cultured S. intermedius are frequently exposed to local unfavorable water temperatures. It has been well documented that S. intermedius are greatly impacted by elevated water temperatures in summer. Besides, acute changes of water temperature also result in negative impacts on regional aquaculture. For example, water temperature frequently decreases rapidly from 22 to 16 °C by the cold water mass at Haiyang island near Dalian (39° 03' N, 123°09' E) in summer, causing mass mortality of cultured species such as the scallop Chlamys farreri. Enhancing resistance abilities can improve the survival of sea urchins at adverse water temperatures. Food is one of the most important methods to facilitate biological resistance abilities. Dietary intervention successfully reduced the mortality of aquacultural Australian greenlip abalone Haliotis laevigata in unfavorable conditions. However, whether S. horneri and S. japonica have potential application for improving the resistance abilities of S. intermedius to adverse water temperatures is not known.

Sea urchins have fitness-related behaviors including righting, tube feet extension and Aristotle's lantern reflex. Righting behavior refers to an inverted sea urchin placed on its aboral surface to correct the posture with the aboral side up and has been commonly using as a stress indicator. Tube feet extension, which refers to the ability of sea urchins to extend their tube feet, is important for the fitness of sea urchins. The Aristotle's lantern reflex is the process of opening and closing of the teeth, which affects the capacity of sea urchins to grasp food with their teeth.

The main purposes of the present study are to investigate: (1) whether S. horneri decreases the mortality and morbidity of sea urchins, compared with S. japonica; (2) whether S. horneri slows the gonadal development of S. intermedius, compared with S. japonica; (3) whether S. horneri and S. japonica have potential application for improving the resistance of S. intermedius under adverse temperatures.

### Results

#### Crude protein, fiber, fat and ash of S. horneri and S. japonica.

The concentrations of crude protein (186.33 ± 2.66 g/kg) and crude fat (10.00 ± 4.36 g/kg) of dried S. horneri were not significantly different from those of dried S. japonica (210.50 ± 6.15 g/kg, t = 1.240, P = 0.283 for crude protein; 6.00 ± 2.83 g/kg, t = 1.315, P = 0.259 for crude fat). However, the concentrations of crude fiber (46.67 ± 6.80 g/kg) and ash (13.94 ± 4.02 g/kg) of dried S. horneri were significantly larger than those of dried S. japonica (31.50 ± 9.19 g/kg, t = 2.910, P = 0.044 for crude fiber; 1.95 ± 0.39 g/kg, t = 5.128, P = 0.007 for ash).

### Experiment I. Number of survivors and diseased sea urchins.

Diet did not significantly affect the number of survivors and diseased sea urchins, compared with S. japonica (χ² = 0.116, P = 1.000, Table 1). However, S. intermedius fed S. horneri showed significantly fewer diseased individuals than those fed S. japonica (χ² = 4.421, P = 0.036, Table 1).

Dried food consumption. The average dried food consumption of S. intermedius fed S. horneri (1.12 ± 0.29 g ind⁻¹ day⁻¹) was significantly more than those fed S. japonica (0.14 ± 0.08 g ind⁻¹ day⁻¹, P < 0.001, Fig. 1).

Test diameter and wet body weight. Compared with the initial conditions, test diameter significantly increased in both treatments (t = 4.272, P < 0.001 for S. intermedius fed S. horneri; t = 5.791, P < 0.001 for S. intermedius fed S. japonica). However, there was no pronounced difference in wet body weight between groups (t = 1.784, P = 0.081 for S. intermedius fed S. horneri; t = 1.002, P = 0.321 for individuals fed S. japonica).

After ~9 weeks, test diameter and wet body weight of S. intermedius fed S. horneri (30.09 ± 2.93 mm for test diameter; 9.86 ± 2.34 g for wet body weight) were not significantly different from those fed S. japonica (30.19 ± 1.87 mm, t = 0.162, P = 0.872 for test diameter; 9.21 ± 1.48 g, t = 1.237, P = 0.223 for wet body weight).

Specific growth rate. There was no significant difference in specific growth rate (0.188 ± 0.189 for S. intermedius fed S. horneri; 0.080 ± 0.192 for individuals fed S. japonica) between both diet treatments (t = 0.891, P = 0.399).

Wet gut weight. No significant difference in wet gut weight was detected between treatments (0.19 ± 0.05 g for S. intermedius fed S. horneri; 0.13 ± 0.05 g for sea urchins fed S. japonica; t = 1.900, P = 0.087).

|          | Sargassum horneri | Saccharina japonica | χ²  | P   |
|----------|------------------|---------------------|-----|-----|
| Survivors| 96               | 95                  | 0.116 | 1.000 |
| Dead     | 4                | 5                   |      |     |
| Diseased | 8                | 18                  | 4.421 | 0.036 |
| Not diseased | 92     | 82                  |      |     |
However, Aristotle's lantern reflex of *Strongylocentrotus intermedius* consistently tube feet extension rating (3.96 ± 1.05) and Aristotle's lantern reflex (4.04 ± 1.87 times min⁻¹) was significantly higher than those fed *Saccharina japonica*.

**Figure 1.** Dried food consumption (g ind⁻¹ day⁻¹) of *Strongylocentrotus intermedius* fed different diets for six consecutive days (mean ± SD, N = 5). *Sargassum horneri* and *Saccharina japonica* refer to the experimental and control groups, respectively. The asterisks *** mean P < 0.001.

**Aristotle’s lantern length and wet Aristotle’s lantern weight.** Aristotle’s lantern length (7.88 ± 0.74 mm) and wet Aristotle’s lantern length (0.46 ± 0.05 g) of *S. intermedius* fed *S. horneri* were not significantly different from those fed *S. japonica* (7.97 ± 0.34 mm, t = 0.275, P = 0.789 for Aristotle’s lantern length; 0.40 ± 0.06 g, t = 2.036, P = 0.069 for wet Aristotle’s lantern weight).

**Gonadal yield.** Compared with the initial conditions, no significant differences were found in wet gonad weight (0.58 ± 0.45 g for *S. intermedius* fed *S. horneri*; 0.63 ± 0.35 g for individuals fed *S. japonica*; t = 0.195, P = 0.850) and gonad index (5.83 ± 1.90 for *S. intermedius* fed *S. horneri*; 7.17 ± 1.40 for individuals fed *S. japonica*; t = 0.564, P = 0.585) in either treatment.

**Gonadal development.** After ~9 weeks, all of the gonads of *S. intermedius* fed *S. horneri* were in the growing stage (stage II). However, 83.33% and 16.67% of individuals fed *S. japonica* were in the premature gonads (stage III) and growing stage (stage II), respectively.

In the gonads of *S. intermedius* fed *S. horneri*, the primary oocytes only attached to the follicular wall of ovaries (Fig. 2A) and sperms only occurred in the follicular wall of testes (Fig. 2B). Regarding the gonads of sea urchins fed *S. japonica* at the same period, the oocytes were detached from the wall and gradually replaced nutritive phagocytes in the follicular cavity (Fig. 2C). Consistently, the basophilic clusters of sperm with a length of ~2 μm occurred in both the follicular wall and follicular cavity of testes (Fig. 2D).

**Experiment II.** **Moderately elevated temperatures.** Significant difference was not found between the treatments in either righting response time (89.3 ± 57.6 s for *S. intermedius* fed *S. horneri*, 74.9 ± 48.5 s for sea urchins fed *S. japonica*, Kruskal–Wallis H = 1.182, P = 0.277; Fig. 3A) or tube feet extension rating (3.32 ± 0.93 for *S. intermedius* fed *S. horneri*, 3.55 ± 0.90 for individuals fed *S. japonica*, Kruskal–Wallis H = 3.814, P = 0.051; Fig. 3B). However, Aristotle’s lantern reflex of *S. intermedius* fed *S. horneri* (4.40 ± 0.95 times min⁻¹) was significantly higher than those fed *S. japonica* (2.49 ± 1.39 times min⁻¹, t = 3.343, P = 0.003, Fig. 3C).

**Acute changes in water temperatures.** The righting response time of *S. intermedius* fed *S. horneri* (78.9 ± 49.9 s) was significantly higher than those fed *S. japonica* (38.6 ± 11.6 s, Kruskal–Wallis H = 17.149, P < 0.001, Fig. 3D). Consistently, tube feet extension rating (3.96 ± 1.05) and Aristotle’s lantern reflex (4.04 ± 1.87 times min⁻¹) of individuals fed *S. japonica* were significantly greater than those fed *S. horneri* (2.94 ± 1.27, Kruskal–Wallis H = 33.872, P < 0.001 for tube feet extension rating, Fig. 3E; 2.44 ± 1.94 times min⁻¹, Kruskal–Wallis H = 6.281, P = 0.012 for Aristotle’s lantern reflex, Fig. 3F).
Discussion

The increased incidence of bacteria causes black-mouth and spotting diseases greatly decreased the production of *S. intermedius* aquaculture\(^1,^4,^8\). We found that *S. intermedius* fed *S. horneri* showed significantly fewer morbidity than those fed *S. japonica*. A reasonable explanation is that polysaccharides enriched in *S. horneri*\(^30\) stimulate the innate immune system of sea urchins because cell walls of pathogenic bacteria possess polysaccharides that are identified as characteristic antigen molecules by the innate immune system\(^31–^33\). This indicates that *S. japonica* could be an effective approach to the disease prevention in *S. intermedius*.

*Strongylocentrotus intermedius* necessarily requires at least one summer to develop from fertilized eggs to adults of commercial size (> 5 cm of test diameter) in long line culture in China\(^4,^7\). It is not essential for small sea urchins to develop gametes, but more appropriate for somatic growth in aquaculture\(^34\). Gonadal precocity greatly consumes the stored energy and consequently leads to poor somatic growth\(^35,^36\) and probable mortality of sea urchins. Dietary protein is the basis of gonadal developments of sea urchins\(^35,^37\). The present study showed that *S. intermedius* fed *S. horneri* exhibited slower gonadal development than those fed *S. japonica*, even though no significant difference of crude protein concentration were found between the two brown algae. This indicates that gonadal development of sea urchins is probably due to other nutrient elements. Collectively, the present study indicates that *S. horneri* is an effective food to avoid the precocious puberty of *S. intermedius* and may subsequently contribute to their energy storage.

In addition, cultured *S. intermedius* requires higher resistance ability at adverse water temperatures in summer. Behaviors, which are realized by the coordination of neuromuscular systems\(^38,^39\), display a strong correlation with the fitness of sea urchins\(^40\). Water temperature significantly affects neuromuscular activities. For example, the sea urchin *Strongylocentrotus purpuratus* showed a decreased adhesion when being exposed to the elevated water temperatures\(^41\). For the first time, the present study found that different species of brown algae have different effects on sea urchin behaviors under acutely changed and moderately elevated temperatures. Specifically, sea urchins fed *S. horneri* showed significantly greater Aristotle’s lantern reflex than those fed *S. japonica* when being exposed to moderately elevated temperatures (an increase from 23.5 to 26.5 °C at a rate of 0.5 °C per day and maintained at 26.5 °C for 1 week). The Aristotle’s lantern reflex represents the ability to operate sea urchin jaws to grasp a food around and is commonly used as an indicator for the food intake capacity\(^29,^42\). Consistently, *S. intermedius* fed *S. horneri* exhibited significantly higher food consumption than those fed *S. japonica*. The study indicates that *S. intermedius* fed *S. horneri* displays a significantly greater capacity in thermal tolerance than those fed *S. japonica*. In comparison, sea urchins fed *S. japonica* outperformed those fed *S. horneri* in behaviors of righting, tube feet extension and Aristotle’s lantern reflex under acutely changed water temperatures. These

---

**Figure 2.** Histology of ovaries (A,C) and testes (B,D) of *Strongylocentrotus intermedius* (N = 6) fed different brown algae after ~ 9 weeks. *Sargassum horneri* and *Saccharina japonica* refer to the experimental and control groups, respectively. In the gonads of sea urchins in the experimental group, the primary oocytes only attached to the follicular wall of ovaries (A) and sperms only occurred in the follicular wall of testes (B). Regarding the gonads of sea urchins in the control group over the same period, the oocytes were detached from the wall and gradually replaced nutritive phagocytes in the follicular cavity (C). Consistently, the basophilic clusters of sperm with a length of ~ 2 μm occurred in both the follicular wall and follicular cavity of testes (D). NP means nutritive phagocytes.
findings clearly suggest that *S. intermedius* fed *S. japonica* is more suitable for the areas with cold water mass in summer because it probably reduces the negative impacts of acute changes in water temperature on sea urchins. A possible explanation is that *S. japonica* contributes to the reduction of the free-radical level in organisms, decreases tissue hypoxia and subsequently improves the neuromuscular activities of sea urchins.

In conclusion, *S. horneri* has direct application potential for the long line culture of *S. intermedius* in summer, because of the advantages in health, energy storage and resistance abilities. Further, *S. japonica* is appropriate for *S. intermedius* aquaculture in the areas with cold water mass where acute changes of water temperature...
exist. The present study provides valuable information into the management of the long line culture of small *S. intermedius* in summer.

**Materials and methods**

**Sea urchins.** Experimental sea urchins were produced in November 2018, fed *Ulva pertusa* ad libitum for ~2 months until the test diameter reached 0.3–0.4 cm diameter and subsequently fed *S. japonica* for culture45. Three hundred healthy *S. intermedius* (~3 cm of test diameter) were randomly selected from an aquaculture farm in Huangnichuan, Dalian (121° 45’ N, 38° 82’ E) and then were transported to the Key Laboratory of Marine Culture and Stock Enhancement in North China’s Sea, Ministry of Agriculture at Dalian Ocean University (121° 56’ N, 38° 87’ E) on 9 July 2019. Sea urchins were maintained in a large fiberglass tank (length × width × height: 180 × 100 × 80 cm) of the recirculating system (Huixin Co., Dalian, China) to acclimatize to laboratory conditions and fed *S. japonica* ad libitum for 1 week with aeration. Water quality parameters were measured daily. Water temperature was 23.55 ± 0.07 °C, pH 7.72 ± 0.02 and salinity 33.76 ± 0.04. They were then fasted for another week until the experiment began.

Test diameter, wet body weight and wet gonad weight were evaluated for the initial conditions of sea urchins before the experiments started (N = 20 for test diameter and wet body weight; N = 10 for wet gonad weight).

**Crude protein, fiber, fat and ash of *S. horneri* and *S. japonica*.** Samples were taken from each dried brown alga to investigate their organic composition (crude protein, crude fiber and crude fat) and ash on 20 August 2019 (N = 3). Semi-micro Kjeldahl nitrogen was used to determine the crude protein concentration of the dried brown algae44. In order to measure the crude fiber concentration of brown algae, about 10 g of each sample of the dried brown alga was boiled with a mixed solution (1.25% dilute acid and dilute alkali) for 30 min and ashed at 550 °C to remove the minerals45. Five grams of each dried sample and 15 mL petroleum ether were added to the Soxhlet extractor and refluxed at constant temperature (45 ± 1 °C) for eight hours to assess the crude fat concentration of the brown algae46. To investigate the ash concentration of the brown algae, approximate two g dried samples were placed in a constant weight fritted glass and burned in a muffle furnace (M110, Thermo CO., U.S) at 550 °C for 48 h46.

**Experiment I.** Experimental design. Diet was the experimental factor, either *S. horneri* or *S. japonica*. Fresh *S. horneri* were collected from a farm in Huangnichuan Dalian (121° 45’ N, 38° 82’ E) and *S. japonica* from Dalian Bay (120° 37’ E 38° 56’ N) in July 2019. Individuals were fed dried *S. horneri* (experimental group) and dried *S. japonica* (control group) ad libitum for ~9 weeks during the experiment (from 23 July 2019 to 25 September 2019). One large fiberglass tank was used for each experimental treatment. One hundred sea urchins were haphazardly chosen and put into 100 individual cylindrical cages (length × width × height: 10 × 10 × 20 cm; 1.5 cm of mesh size) in each tank (length × width × height: 150 × 100 × 60 cm) of the recirculating system (Huixin Co., Dalian, China) with aeration, according to the experimental design. Diseased sea urchins were removed timely from the tanks to avoid the potential spread of infectious diseases in experimental treatments and were transported into new tanks (length × width × height: 75 × 45 × 35 cm) for individual culture and observation following with the previous management.

Water temperature was not controlled, ranging from 21.3 to 25.6 °C during the experiment. Water quality parameters were measured weekly as pH 7.59–7.85 and salinity 32.69–32.13. One-half of the seawater was renewed daily.

**Number of survived and diseased sea urchins.** Black-mouth disease refers to the perioral membrane turns black (Fig. 4A) with the decreased ability of attaching and feeding in sea urchins47. Sea urchin with spotting disease is indicated by the spotting lesions with red, purple or blackish color on the body wall followed by the detachment of local spines48 (Fig. 4B). The enlarging spotting lesions commonly cause ulceration on the body wall and finally result in death8. Sea urchin without disease performance is shown in Fig. 4C. The number of survived and diseased sea urchins (either black-mouth or spotting diseased) was recorded during the experiment.

**Dried food consumption.** The measurement of food consumption was conducted for six consecutive days (from 7 August 2019 to 12 August 2019). The total supplemented and remained diets were weighed (G & G Co., San Diego, USA) after removing the water on their surface. The samples of uneaten diets were collected, weighed and dried for 4 days at 80 °C and then reweighed (N = 5). To avoid the loss of uneaten food, a fine silk net (mesh size 260 μm) was set outside the cage to collect the fragments of uneaten brown algae.

Dried food consumption was calculated as follows (according to Zhao et al.49 with some revisions):

\[
F = W_1 \times (1 - \frac{B_s - B_u}{B_s}) - W_2 \times (1 - \frac{C_s - C_u}{C_s})
\]

F = dried food consumption (g), \(W_1\) = wet weight of total supplement diets (g), \(W_2\) = wet weight of total uneaten diets (g), \(B_s\) = wet weight of sample supplemented diets (g), \(B_u\) = dry weight of sample supplemented diets (g), \(C_s\) = wet weight of sample uneaten diets (g), \(C_u\) = dry weight of sample uneaten diets (g).

**Growth.** Test diameters, Aristotle’s lantern length were measured using a digital vernier caliper (Mahr Co., Ruhr, Germany). Body, Aristotle’s lantern and gut were weighed wet using an electric balance (G & G Co., San Diego, USA) on 25 September 2019 (N = 27 for test diameter and wet body weight; N = 6 for Aristotle’s lantern length, wet weight of Aristotle’s lantern and gut).
Specific growth rate (SGR) was calculated according to the following formula:

$$\text{SGR} (%) = \left( \frac{\ln P_2 - \ln P_1}{D} \right) \times 100$$

SGR = specific growth rate, $P_2$ = final wet body weight, $P_1$ = initial wet body weight, $D$ = experimental duration.

Gonad yield. Gonads were carefully collected from each treatment and weighed using an electric balance (G & G Co., San Diego, USA) on 25 September 2019 (N = 6). Gonad index was calculated according to the following formula:

$$\text{GI} (%) = \left( \frac{GW}{BW} \right) \times 100$$

GI = gonad index, GW = wet gonad weight, BW = wet body weight.

Gonadal development. One of five pieces of each gonad was preserved in the Bouin’s solution (saturated picric acid solution: formaldehyde: glacial acetic acid = 15: 5: 1) for 48 h between the treatments (N = 6). Standard histology technique, including embedment, infiltration, section and stain, was performed to make the gonad tissue slices. Sections were classified according to the stage of development of germinal cells and nutritive phagocytes: stage I, recovering; stage II, growing; stage III, premature; stage IV, mature; stage V, partly spawned; stage VI, spent.

Experiment II. Experimental design. Experiment II lasted for 4 weeks (from 25 September 2019 to 23 October 2019). Eighty healthy sea urchins were haphazardly selected from each treatment at the end of experiment I. They were then distributed into 80 cylindrical cages (5×10×10 cm) in each fiberglass tanks (length × width × height: 77.5×47.0×37.5 cm) of the temperature-controlled system (Huixin Co., Dalian, China) with aeration in both treatments. Sea urchins were maintained at 23.5 °C for 2 weeks (the average water temperature of experiment I) to eliminate the past thermal history, following the previous diet strategy of experiment I. Water quality was recorded daily as pH 7.83–7.85 and salinity 32.44–32.62. One-third of the seawater was renewed daily.

**Figure 4.** The conceptual diagrams show the black-mouth disease (A), spotting disease (B) and without disease performance (C) of sea urchin as well as the devices for righting behavior (D) and Aristotle’s lantern reflex (E).
Subsequently, to investigate whether *S. horneri* and *S. japonica* contribute to the resistance abilities of small *S. intermedius* at moderately elevated temperatures, 40 individuals were haphazardly chosen from each treatment and placed into 40 cylindrical cages (5 × 10 × 10 cm) in each tank (length × width × height: 77.5 × 47.0 × 37.5 cm) of a temperature-controlled system (Huixin Co., Dalian, China) with aeration for both groups on 9 October 2019. They were subsequently exposed to the moderately elevated temperatures (rose from 23.5 to 26.5 °C at a rate of 0.5 °C per day and maintained at 26.5 °C for 1 week), according to the records of water temperature in Heishijiao sea area (~ 2 m water depth, 38° 51′ N, 121° 33′ E) in the summer of 2017 and 2018 (Fig. 5). Righting behavior, tube feet extension and Aristotle’s lantern reflex were assessed on 23 October 2019.

Similarly, to explore the effects of *S. horneri* and *S. japonica* on the resistance abilities of small *S. intermedius* under acute changes of water temperature, another 40 individuals were randomly selected and placed into 40 cylindrical cages (5 × 10 × 10 cm) in each tank (length × width × height: 77.5 × 47.0 × 37.5 cm) of the temperature-controlled system (Huixin Co., Dalian, China) with aeration for both treatments on 9 October 2019. The water temperature was set at 23.5 °C. A tank of seawater was prepared at 15 °C. To simulate the changes of water temperature in Haiyang island near Dalian (39° 03′ N, 123° 09′ E) where water temperature frequently fluctuates from 22 to 16 °C instantly by the cold water mass17, sea urchins were transferred directly from 23.5 to 15 °C, maintained at 15 °C for an hour and subsequently quickly returned to 23.5 °C for another hour to finish one cycle of the acute change of water temperature. After four cycles, righting behavior, tube feet extension and Aristotle’s lantern reflex of sea urchins were observed.

**Righting behavior.** Sea urchins were placed with the aboral side down on the bottom of an experimental tank (length × width × height: 60 × 40 × 16 cm, Fig. 4D). Righting response time is the time required for individuals in the inverted posture to right themselves with the aboral side up22. The righting response time in seconds was recorded during 10 min. If individuals did not right themselves within 10 min, the time was counted as 600 s (N = 15).

**Tube feet extension.** The method of assessing tube feet extension was established according to You et al.27, with some revisions. Sea urchins were maintained in a tank (length × width × height: 12 × 10 × 10 cm) with fresh seawater for ~ 5 min before the observation (N = 15). The subjective assessment of tube feet extension was evaluated by a well-trained team (5 persons) that was familiar with tube feet extension analysis of sea urchins. The ranking method was quantified based on the quantity and length of tube foot.

Tube feet extension (rating 1–5):

1 = extremely abnormal (not extending)
2 = severe abnormality (extremely low quantity and extremely short length)
3 = moderate anomaly (low quantity and short length)
4 = mild abnormality (slight decrease in quantity and length)
5 = normal (normal quantity and length)
Aristotle’s lantern reflex. A simple device, which has two small compartments (length × width × height: 4.8 × 5.6 × 4.5 cm) with a food film on the bottom, was used to measure Aristotle’s lantern reflex according to our previous study29. Food film was made by a mixture of ~2.5 g agar and 50 ml seawater in order to avoid the potential impacts of the food palatability on sea urchins. The number of Aristotle’s lantern reflex were counted within 5 min using a digital camera (Canon Co., Shenzhen, China) under the device (N = 7 for sea urchins fed S. horneri and N = 10 for individuals fed S. japonicus under moderately elevated temperatures; Fig. 4E).

Statistical analysis. Normal distribution and homogeneity of variance of the data were analyzed using the Kolmogorov–Smirnov test and Levene test, respectively. The number of survived and diseased S. intermedius were compared using the Fisher’s exact test. Food consumption was analyzed using one-way repeated measured ANOVA. Kruskal–Wallis test was performed to compare the difference between the final and initial conditions of sea urchins. Test diameter, wet body weight, SGR, Aristotle’s lantern length, wet Aristotle’s lantern weight, gonad index and Aristotle’s lantern reflex (under moderately elevated temperatures) were analyzed using the independent-samples t test. All data analyses were performed using SPSS 19.0 statistical software. A probability level of P < 0.05 was considered significant.

Received: 30 March 2020; Accepted: 10 July 2020
Published online: 27 July 2020

References
1. Chang, Y., Ding, J., Song, J. & Yang, W. Biology and Aquaculture of Sea Cucumbers and Sea Urchins (Ocean Press, Beijing, 2004).
2. Unuma, T. Introduction: Sea urchin fisheries in Japan. In Echinoderm Aquaculture (eds Brown, N. P. & Eddy, S. D.) 77–85 (Wiley, Hoboken, 2015).
3. Agatsuma, Y. Stronglylocentrotus intermedius. In Developments in Aquaculture and Fisheries Science (ed. Lawrence, I. M.) (Elsevier, Amsterdam, 2013).
4. Lawrence, J. M., Zhao, C. & Chang, Y. Q. Large-scale production of sea urchin (Strongylocentrotus intermedius) seed in a hatchery in China. Aquacult. Int. 27, 1–7. https://doi.org/10.1007/s1049-018-0319-2 (2019).
5. Zhang, X. China Fishery Statistical Yearbook 2019 (China Agriculture Press, Beijing, 2019).
6. Chang, Y. Q. et al. Aquaculture of the sea urchins Stronglylocentrotus intermedius in Fujian coastal areas. South China Fish. Sci. 16, 1–9. https://doi.org/10.12131/210910156 (2019).
7. Zhang, L. et al. Gulfweed Sargassum horneri is an alternative diet for aquaculture of juvenile sea urchins Stronglylocentrotus intermedius in summer. Aquacult. Int. 25, 905–914. https://doi.org/10.1007/s1049-016-0088-8 (2017).
8. Zhang, W. et al. Transcriptome profiling reveals key roles of phagosome and NOD-like receptor pathway in spotted disease Stronglylocentrotus intermedius. Fish Shellfish Immunol. 84, 521–531. https://doi.org/10.1016/j.fsi.2018.04.012 (2019).
9. Lin, Q., Wu, J. S. & Zeng, Z. N. Comparative study on culture biology of triploid and diploid induced by Cyclosporin a in sea urchin Strongylocentrotus horneri (Edwards, 1874). Fish Sci. Technol. Inf. 28, 265–267 (2001) [in Chinese].
10. Kong, Y. T., Cheng, Z. M., Wang, Q. & Zhong, W. Annual reproductive cycle of sea urchin, Stronglylocentrotus intermedius, in raft cultivation. Fish. Sci. 21, 18–21 (2002) [in Chinese].
11. Joll, L. & Caputi, N. Geographic variation in the annual reproductive cycle of the saucer scallop, Amusium ballotii, (Bernardi, 1861) (Molusca: Pectinidae), along the Western Australian coast. Mar. Freshw. Res. 46, 779. https://doi.org/10.1071/MF9950779 (1995).
12. Apraiz, I., Mi, J. & Cristobal, S. Identification of proteomic signatures of exposure to marine pollutants in Mussels (Mytilus edulis). Mol. Cell. Proteomics 5, 1274–1285. https://doi.org/10.1074/mcp.M500333-MCP2006 (2006).
13. Hokkaido Central Fisheries Experimental Station, Hiribeshiboku Fisheries Extension Office & Hokkaido Institute of Marine Science. On the nature seeds collection, intermediate culture and release of the sea urchin, Stronglylocentrotus intermedius. J. Hokkaido Fish. Exp. Station 41, 270–315 (1984) [in Japanese].
14. Du, B., Zhang, Y. J., Zhang, Y. C. & Wang, H. The characteristic of cold water mass variation at the bottom of the north yellow sea and its hydrological effects on the mortality of shellfish cultured in the water of outer China-Shan islands. Mar. Sci. Bull. 14, 17–28 (1996) [in Chinese].
15. Guo, J., Ji, D., Hou, C., Guo, K. & Ji, L. Impact of tropical cyclone Matmo on mixed zone of the Yellow and Bohai Seas. J. Oceanol. Limnol. 36, 1484–1493. https://doi.org/10.31433/10-0785-x (2018).
16. Zhou, W. Water temperature characteristics of Oceanic islands which were affected by the yellow sea cold water mass. Acta Oceanol. Sin. 18, 113–118 (1996) [in Chinese].
17. Zhou, W. et al. The relation between the abnormal change of water temperature and the scallop (Chlamys farreri) death in Haiyang island sea area. Trans. Oceanol. Limnol. 4, 56–62 (1992) [in Chinese].
18. Sangha, J. S. et al. Bioactive components of the edible strain of red alga, Chondrus crispus, enhance oxidative stress tolerance in Caenorhabditis elegans. J. Funct. Foods 6, 1180–1190. https://doi.org/10.1016/j.jff.2013.04.001 (2013).
19. Cheng, M. et al. Antioxidant activity of the polysaccharide fraction of a marine chlorophycean macroalga in Oreochromis niloticus (Linnaeus, 1758). J. Appl. Ichthyol. 34, 556–567. https://doi.org/10.1111/jai.13606 (2018).
20. Stone, D. A. J. et al. Dietary intervention improves the survival of cultured greenlip abalone (Haliotis laevigata Donovan) at high water temperature. Aquaculture 430, 230–240. https://doi.org/10.1016/j.aquaculture.2014.03.047 (2014).
21. Hyman, L. H. The Invertebrates: Echinodermata (McGraw-Hill, New York, 1955).
22. Chi, X. et al. Fitness benefits and costs of shelters to the sea urchin Glyptochorioides crenulatus. PeerJ 8, e8886. https://doi.org/10.7717/ peerj.8886 (2020).
23. Lawrence, J. M. The effect of temperature-salinity combinations on the functional well-being of adult Lytechinus variegatus (Lamarck) (Echinodermata, Echinolidae). J. Exp. Mar. Biol. Ecol. 18, 271–275. https://doi.org/10.1016/0022-0981(75)90111-2 (1975).
24. Hagen, N. T. Is righting response a useful indicator of functional well-being in the green sea urchin Stronglylocentrotus droebachiensis. In Echinoderms Through Time (eds David, G. & Redi, E.) 693–698 (Balkema Press, Rotterdam 1994).
25. Böttger, S. A., McClintock, J. B. & Klinger, T. S. Effects of inorganic and organic phosphates on feeding, feeding absorption, nutrient allocation, growth and righting responses of the sea urchin Lytechinus variegatus. Mar. Biol. 138, 741–751. https://doi.org/10.1007/s00227-006-0476 (2001).
26. Santos, R. & Flammang, P. Intra- and interspecific variation of attachment strength in sea urchins. *Mar. Ecol. Prog. Ser.* **332**, 129–142. https://doi.org/10.3354/meps332129 (2007).

27. You, K., Zeng, X. Q., Liu, H., Zhang, X. M. & Liu, Q. Selectivity and tolerance of sea urchins (*Hemicentrotus pulcherrimus*) to environmental change. *Chin. J. Appl. Ecol.* **14**, 409–412 (2013) (in Chinese).

28. Brothers, C. J. & McClintock, J. B. The effects of climate-induced elevated seawater temperature on the covering behavior, righting response, and Aristotle’s lantern reflex of the sea urchin *Lytechinus variegatus*. *J. Exp. Mar. Biol. Ecol.* **467**, 33–38. https://doi.org/10.1016/j.jembe.2015.02.019 (2015).

29. Ding, J. *et al.* Effects of water temperature on survival, behaviors and growth of the sea urchin *Mesocentrotus nudus*: New insights into the stock enhancement. *Aquaculture* **519**, 734873. https://doi.org/10.1016/j.aquaculture.2019.734873 (2020).

30. Wen, Z. S. *et al.* Composition and anti-inflammatory effect of polysaccharides from *Sargassum horneri* in RAW264.7 macrophages. *Int. J. Biol. Macromol.* **88**, 403–413. https://doi.org/10.1016/j.ijbiomac.2016.02.025 (2016).

31. Urbe, C., Polch, H., Enriquez, R. & Moran, G. Innate and adaptive immunity in teleost fish: A review. *Vet. Med.* **56**, 486–503. https://doi.org/10.17221/3294-VETMED (2011).

32. Skjermo, J. *et al.* Immunostimulation of juvenile turbot (*Scophthalmus maximus L.*) using an alginate with high mannuronic acid content administered via the live food organism *Artemia.* *Fish Shellfish Immunol.* **5**, 531–534. https://doi.org/10.1016/s1000-4648(95)80053-0 (1995).

33. Cheng, A., Tu, C., Chen, Y., Nan, F. & Chen, J. The immunostimulatory effects of sodium alginate and iota-carrageenan on orange-spotted grouper *Epinephelus coioides* and its resistance against *Vibrio alginolyticus.* *Fish Shellfish Immunol.* **22**, 197–205. https://doi.org/10.1016/j.fsi.2006.04.009 (2007).

34. Lawrence, J. M. *Conflict Between Somatic and Gonadal Growth in Sea Urchins: A Review* (2000). https://crdpm.umcs.ca/OURSI/N.

35. Tianyang Zhang, Jiangnan Sun, Xiyuan Huang and Huiyan Wang for their assistance.

36. Skjermo, J. *et al.* Immunostimulation of juvenile turbot (*Scophthalmus maximus L.*) using an alginate with high mannuronic acid content administered via the live food organism *Artemia.* *Fish Shellfish Immunol.* **5**, 531–534. https://doi.org/10.1016/s1000-4648(95)80053-0 (1995).

37. Kelly, M. S., Brodie, C. & McKenzie, J. D. Somatic and gonadal growth of the sea urchin, *Psammechinus miliaris*, maintained in polyculture with the Atlantic salmon. *J. Shellfish Res.* **17**, 1557–1562 (1998).

38. De Ridder, C. & Lawrence, J. M. Food and feeding mechanisms: Echinoida. In *Echinoderm Nutrition* (eds Jangoux, M. & Lawrence, M.) 57–115 (Balkema, Rotterdam, 1982).

39. Kleitman, N. The effect of temperature on the righting of Echinodermata. *Biol. Bull.**80**, 292–298. https://doi.org/10.2307/15377716 (1941).

40. Ling, S. D. & Johnson, C. R. Marine reserves reduce risk of climate-driven phase shift by reinstating size- and habitat-specific trophic interactions. *Ecol. Appl.* **22**, 1232–1245. https://doi.org/10.1890/11-1587.1 (2012).

41. Percy, J. A. Thermal adaptation in the boreo-arctic echinoid, Strongylocentrotus droebachiensis (O. F. Muller, 1776). III. Seasonal acclimatization and metabolism of tissues in vitro. *Physiol. Zool.* **47**, 59–67. https://doi.org/10.1086/physzool.47.1.30155822 (1974).

42. Agustina, Y. Effect of the covering behavior of the juvenile sea urchin *Strongylocentrotus intermedius* on predation by the spider crab *Pugettia quadriceps*. *Fish. Sci.* **67**, 1181–1183. https://doi.org/10.1006/fsis.2001.06399. (2001).

43. Sprygin, V. G., Kushnerova, N. F., Fomenko, S. E., Sizova, L. A. & Motom, T. V. The hepatoprotective properties of an extract from the brown alga *Saccharina japonica*. *Russ. J. Mar. Biol.* **39**, 65–69. https://doi.org/10.1134/S1063074013010100 (2013).

44. Chen, Z. H., Shi, M., Wang, Q. X. & Zhang, X. H. Protein content measurement of food using the method of Kjeldahl determination. *Xinjiang Anim. Husk.* **5**, 22–24 (2008) (in Chinese).

45. Cheng, J. *Improvement of Determination Method of Feed Conventional Analysis* (Northeast Agricultural University, Harbin, 2016).

46. Zang, J., Yang, J. & Dong, W. Study on improved methods of determination of crude fat in foods. In *Modern Agricultural Science Technology* 333–334 (2012) (in Chinese).

47. Li, T. W., Xu, S. L., Wang, R. B., Xu, S. F. & Su, X. R. Preliminary studies on the black mouth disease of sea urchin, *Strongylocentrotus intermedius*. *Mar. Sci.* **24**, 41–43 (2000) (in Chinese).

48. Wang, Y. N., Chang, Y. Q. & Lawrence, J. M. Sea Urchins: Biology and Ecology. *Disease in Sea Urchins* (Academic Press, Cambridge, 2013).

49. Zhao, C. *et al.* Effects of temperature and feeding regime on food consumption, growth, gonad production and quality of the sea urchin *Strongylocentrotus intermedius*. *J. Mar. Biol. Assoc. U.K.* **96**, 185–195. https://doi.org/10.1017/S0025384115001617 (2016).

50. Johnstone, J., Nash, S., Hernandez, E. & Rahman, M. S. Effects of elevated temperature on gonadal functions, cellular apoptosis, and oxidative stress in Atlantic sea urchin *Arbacia punctulata*. *Mar. Environ. Res.* **149**, 40–49. https://doi.org/10.1016/j.marenvres.2019.05.017 (2019).

51. Byrne, M. Annual reproductive cycles of the commercial sea urchin *Paracentrotus lividus* from an exposed intertidal and a sheltered subtidal habitat on the west coast of Ireland. *Mar. Biol.* **104**, 275–289. https://doi.org/10.1007/BF01313269 (1990).

52. Laegsgaard, P., Byrne, M. & Anderson, D. T. Reproduction of sympatric populations of *Helicolaridis erythrogramma* and *H. tuberculata* (*Echinoida*) in New South Wales. *Mar. Biol.* **110**, 359–374. https://doi.org/10.1007/bf01343535 (1991).

53. King, C. K., Hoegh-Guldberg, O. & Byrne, M. Reproductive-cycle of *Centrostephanus rodgersii* echinoida, with recommendations for the establishment of a sea-urchin fishery in new-south-wales. *Mar. Biol.* **120**, 95–106 (1994).

**Acknowledgements**

This work was supported by Chinese Outstanding Talents in Agricultural Sciences (for Yaqing Chang), Laboratory for Marine Fisheries Science and Food Production Processes, Qingdao National Laboratory for Marine Science and Technology, China P. R. (2017-1B05), a grant for innovative talents in universities in Liaoning Province (for Chong Zhao). All applicable international, national, and/or institutional guidelines for the care and use of animals were followed by the authors. We thank Prof. John Lawrence for academic and editorial suggestions and Tianyang Zhang, Jiangnan Sun, Xiuyan Huang and Huiyan Wang for their assistance.

**Author contributions**

C.Z., F.H. and Y.C. designed the experiments. F.H., M.Y., P.D., X.Z., Z.C. and J.L. carried out the experiments. F.H. and J.D. and X.C. did the data analysis. F.H. and C.Z. wrote the manuscript. All authors gave final approval for publication.

**Competing interests**

The authors declare no competing interests.
