



Larval and Juvenile Development of the Echinometrid Sea Urchin *Colobocentrotus mertensii*: Emergence of the Peculiar Form of Spines

Authors: Thet, May Maw, Noguchi, Masashi, and Yazaki, Ikuko

Source: Zoological Science, 21(3) : 265-274

Published By: Zoological Society of Japan

URL: <https://doi.org/10.2108/zsj.21.265>

Larval and Juvenile Development of the Echinometrid Sea Urchin *Colobocentrotus mertensii*: Emergence of the Peculiar Form of Spines

May Maw Thet^{1*}, Masashi Noguchi² and Ikuko Yazaki¹

¹*Department of Biological Sciences, Graduate School of Science, Tokyo Metropolitan University, 1-1, Minami-oshawa, Hachioji-shi, Tokyo 192-0397, Japan*

²*3-8-5-806, Kitamachi, Nerima-ku, Tokyo 179-0081, Japan*

ABSTRACT—The development of *Colobocentrotus mertensii* from embryos to larvae and early juveniles was observed to give the first detailed description of larval and juvenile formation and skeletal structures in echinometrid sea urchins. The first larval spicules appeared at the mesenchyme blastula stage, whereas, in many echinoids, spicules were formed after gastrulation. From late eight-armed larva to juvenile, body color of *C. mertensii* was deep red, which has never been described for any echinoid before.

The adult form of *C. mertensii* is characteristic in that the spines at the aboral side are short, truncated and pavement-like. The first sign of peculiar adult features could be seen in the juvenile spines and adult spines, which are broader than those of closely related *Anthocidaris crassispina*. The primary podia emerged on the left side of larval body were more stout and thicker in *C. mertensii* than in *A. crassispina*. The present study shows that developmental process of larval structure of *C. mertensii* is in general similar to the *A. crassispina* and the differences is first seen in juvenile structure including the distribution of pigment spots and morphology of adult spine.

Key words: echinometrid, larva, juvenile, spicule, spine

INTRODUCTION

Living echinoids are subdivided into two groups, regular and irregular (Hyman, 1955). *Colobocentrotus mertensii*, an endemic Japanese sea urchin species, is closely related to *Anthocidaris crassispina*, occurs in the warm current from

the Miyake Island in Izu, Ogasawara Islands and the Kii Peninsula to RyuKyu Islands (Shigei, 1974), and is classified as a regular urchin. The test of regular urchins is characterized by its globose shape and long needle-like spines (Hyman, 1955). However, *C. mertensii* is different from other regular sea urchins in that it has a hat-shaped test. More-

* Corresponding author: Tel. +81-426-77-2562;
FAX. +81-426-77-2559.
E-mail: maymaw-thet@c.metro-u.ac.jp

Abbreviations used in figures.

a.ci.b=	anterior ciliated band	j.sp=	juvenile spine	pr.ci.b=	preoral ciliated band
ala=	anterolateral arm	l.ala=	left anterolateral arm	proa=	preoral arm
a.sp=	adult spine	l.alr=	left anterolateral rod	ptr=	posterior transverse rod
am.a=	ambulacral area	l.pcv=	left posterior coelomic vesicle	r.ala=	right anterolateral arm
ap=	anal plate	l.pda=	left posterodorsal arm	r.pda=	right posterodorsal arm
ax=	axocoel	l.pdr=	left posterodorsal rod	r.pdr=	right posterodorsal rod
b=	bud of preoral arm	l.poa=	left postoral arm	r.poa=	right postoral arm
da=	dorsal arch	l.por=	left postoral rod	r.por=	right postoral rod
d.p.p=	disc of primary podium	l.proa=	left preoral arm	r.proa=	right preoral arm
e=	esophagus	l.pror=	left preoral rod	ru.ped=	rudiment of pedicellaria
G=	genital plate	m=	mouth	s=	stomach
f.p=	fenestrated plate	p.ci.b=	posterior ciliated band	sp=	spine
g.jp=	genital juvenile spine	ped=	pedicellaria	T=	terminal plate
hy=	hydrocoel	pig=	pigment spots	trs=	triradiate spicule
i.am.a=	interambulacral area	p.p=	primary podium	v=	vestibule

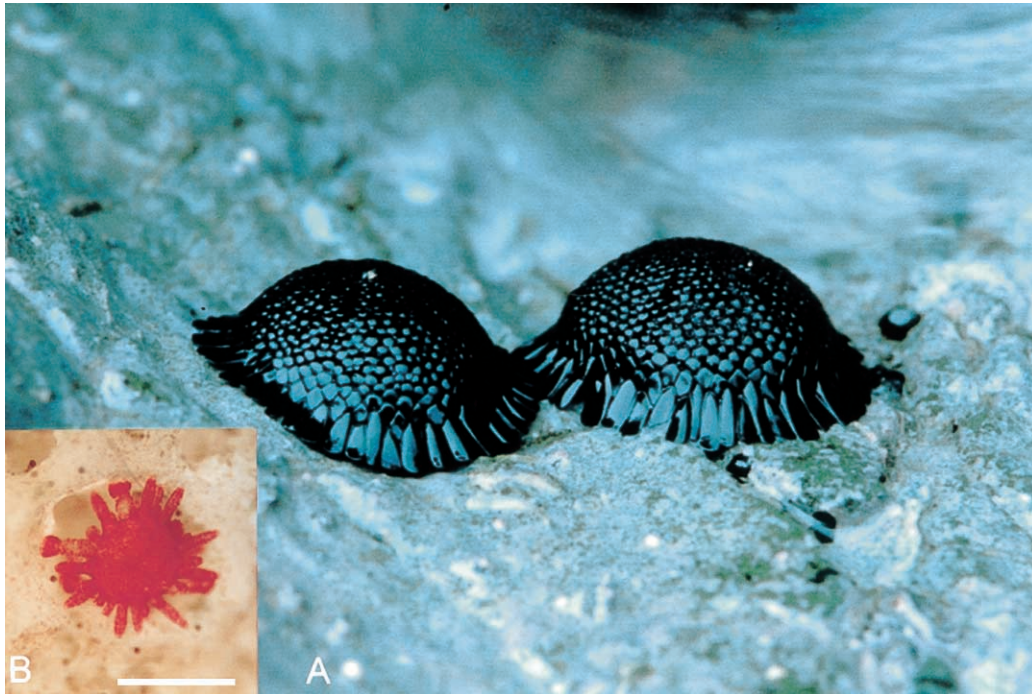


Fig. 1. Two adult individuals (**A**) and a juvenile (**B**) of *Colobocentrotus mertensii*, from Bonin Islands. Adults are olive green in color and have an oval shape. The spines of aboral side are short while the spines of margin are broad, flat and chisel-shaped. The juvenile shows characteristic color due to pigments. Aboral side views. Scale bar in B, 0.5 mm.

over, the spines of the aboral side is very short, truncate, and form pavement-like polygonal plates covering the whole aboral side. The marginal spines are broad, flat and chisel-shaped (Fig. 1A). On the oral side, the tube-feet are well developed, and covered by short and flat spines. Thus *C. mertensii* has an unusual morphology and peculiar spines in adulthood, but it is classified in the family of Echinometridae (Shigei, 1986). In the echinometrid sea urchins, spines of adults are usually needle-like, such as seen in *A. crassispina*, while those of *C. mertensii* are flat in shape.

The aim of the present study is to observe the development of *C. mertensii* to confirm that its morphology in the early stages is similar to those of other members of the family Echinometridae. Here we report that the difference in morphology of *C. mertensii* and *A. crassispina* is first seen in larval and juvenile stages.

As for the development of sea urchins of genus *Colobocentrotus*, Mortensen (1921) reported the development of early stages up to four-armed pluteus of *C. atratus* in Hawaii. There has been, however, no available information about *C. mertensii*. In the case of *A. crassispina*, Onoda (1931) described its development from early stages up to juvenile.

The observations made in the present study demonstrated that the peculiar form of adult spine in *C. mertensii* emerges first in juvenile stages and that the comparison of its developmental morphology with those of other Echinometridae members clearly places *C. mertensii* among Echinometridae.

MATERIAL AND METHOD

Adult specimens of *C. mertensii* were collected from Chichijima Island, the Ogasawara (Bonin) Islands. *A. crassispina* and *Pseudocentrotus depressus* from Bousou Peninsula. Urchins were induced to spawn by intracoelomic injection of 0.5 to 1.0 ml of 0.5 M KCl. The natural sea water was used for culture after filtration and sterilization at 80°C for 20 min. Eggs were washed three times in sterilized sea water and fertilized with a diluted sperm solution. Larvae were cultured with constant stirring of sea water by a paddle at a speed of 30 r.p.m in a 3L glass beaker. They were kept at 27°C for *C. mertensii* and 23°C for *A. crassispina*. After larvae reached early four-armed stage, they were fed diatome *Chaetoceros gracilis* cultivated in the laboratory with medium KW-21, every day or every other day. In this paper, Mortensen's (1921) terminology was used for larval arms and spicules. Observations of metamorphosis were made on late larvae with well-developed primary podia and juvenile spines. Metamorphosing larvae were put into glass bowls with approximately 150 ml of filtered sea water. Each bowl contained a small rock (4–5 cm³) which had been collected from adult habitats and held in the laboratory sea water table prior to use.

Preparation of the skeletal specimens

To make the preparations for drawing or photography, larvae were fixed on the depression slide with a drop of 5% formaline solution in the sea water. Preparation of the larval skeleton and the corona of juvenile was done according to Goldern (1926). The materials fixed in 70% ethanol or living larvae in sea water were washed in the depression slides with distilled water to remove alcohol or salts. Next, the muscular parts of the materials were macerated with 1M KOH and the specimens were rinsed thoroughly in distilled water to remove KOH. They were then washed three times with distilled water and, in order to make the specimens transparent, put in a 50% glycerine in water. Finally, a drop of pure glycerine

was put on the skeletal specimens and slides were covered with a cover glass. Two polarizing plates were used to distinguish overlapped skeletons separately by their respective birefringence. Drawing were made with Nikon's drawing apparatus.

RESULTS

The average developmental time course of *C. mertensii* reared at 27°C is summarized in Table 1. In general, the variation in developmental stages among cultures increased with time, although larvae appeared healthy throughout the culture period. For example, after 37 days,

when the first larvae metamorphosed completely into juveniles, others had only partially developed juvenile structures while still others showed no external sign of juvenile rudiment development.

Early larval development

The mature egg of *C. mertensii* was 69 µm in diameter and slightly yellowish in color and quite opaque. Early cleavage and development followed the pattern similar to those reported for *A. crassispina* (Onoda, 1931). It is to be noted that the appearance of first spicules of *C. mertensii* and *A. crassispina* was earlier than that of non-Echinometridae sea

Table 1. Summary of developmental stages of *Colobocentrotus mertensii*.

Time after fertilization	Developmental stage
2.75 hr	Eight cell embryo.
3.5 hr	Sixteen cell embryo.
12 hr	Mesenchyme blastula. Triradiate spicules evident.
13.5 hr	Gastrula. Archenteron began to invaginate into blastocoel.
17 hr	Late gastrula. The archenteron touched the prospective oral part of the ectoderm.
28 hr	Early two-armed larva. A pair of coelomic vesicle developed on the top of archenteron.
32 hr	Two-armed larva. Fenestrated postoral arms were formed and red pigment spots were scattered through the larval arms.
3 d	Early four-armed larva. After anterolateral arms well developed, larva was fed on <i>C. gracilis</i> .
7 d	Late four-armed larva. Postoral arms well developed, 630 µm long (measured from posterior curve of the larval body to arm tip).
16 d	Posterodorsal arms were about one-third shorter than postoral arms. Buds of preoral arms present.
23 d	Echinus rudiment began to form. Preoral arm developed from dorsal arch. Posterior transverse spicule present.
32 d	Three ciliated bands developed. Postoral arms approx. 800 µm long. The more advanced larva had five primary podia and three pedicellariae and juvenile spines.
35 d	The most advanced larva showed reduction in arm length.
37 d	First larva metamorphosed and the juvenile became deep red. Juvenile test diameter approx. 770 µm including the spine.

Larvae were reared at 27°C.

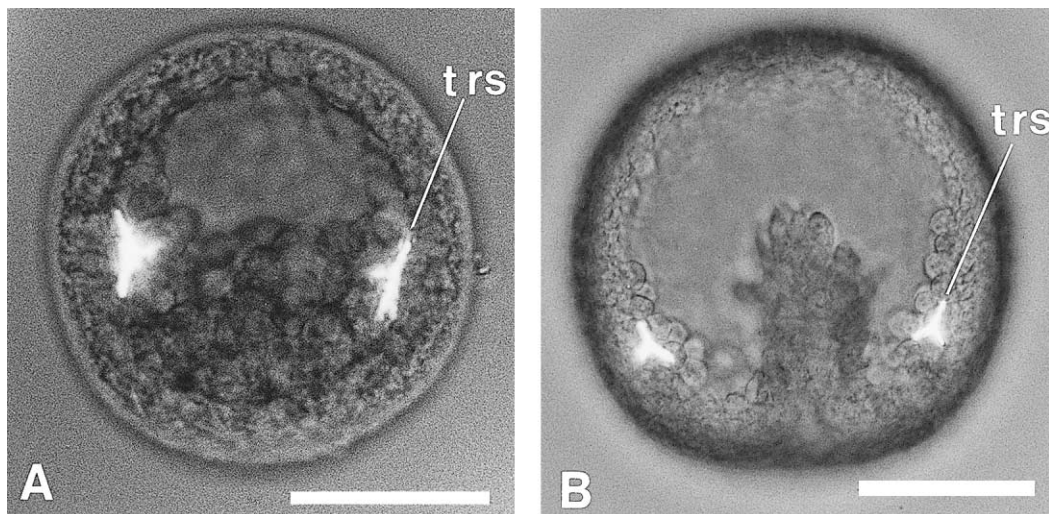


Fig. 2. Birefringence of spicules of mesenchyme blastula stage embryo of *Colobocentrotus mertensii* (A) and mid gastrula stage embryo of *Pseudocentrotus depressus* (B) photographed with polarization microscope. Note that first pair of triradiate spicules appears much earlier in *C. mertensii* than in *P. depressus*. Scale bars in A and B, 50 µm.

urchin species. In *C. mertensii* triradiate spicules (trs) were formed in mesenchyme blastula stage before gastrulation (Fig. 2A) while in many sea urchin species, spicules were formed at mid gastrula stage (Fig. 2B, *Pseudocentrotus depressus*).

At 42 hr, in addition to two fenestrated postoral arms (poa), two anterolateral arms (ala) having simple rod appeared, forming the four-armed echinopluteus. The preoral ciliated band (pr.ci.b) developed in front of the mouth. Red pigment spots (pig) which were initially present throughout the larval arms became increasingly concentrated at the tips after this stage. The four-armed stage persisted through the 7th day after fertilization (Fig. 3A).

Seven to 9 days after fertilization, bases of postoral arms bulged and a triradiate spicule, a rudiment of posterodorsal rod, was formed in each bulge. Soon later, the 3rd triradiate spicule, a rudiment of dorsal arch, was formed. The fenestrated posterodorsal arms (pda) characteristic of the six-armed stage appeared 12 days after fertilization and fully developed by the 16th day (Fig. 3B).

Late larval development

Between the 18th and 20th day, the posterior area of the larval body gradually became flat and dorsal arch (da) elongated anteriorly and formed two preoral arms (proa) indicating the initiation of the eight-armed stage. With further development, a pair of anterior ciliated band (a.ci.b) (vibratile type) grew on the dorsal and ventral surface (Fig. 4A). During the late six-armed stage, the left and right coelomic

vesicles were divided into anterior and posterior parts. Left anterior coelomic vesicle formed a hydrocoel (hy), a rudiment of adult water-vascular system (MacBride, 1903, 1914; Fukushi, 1960), and an axocoel (ax). Twenty three days after fertilization, the ectodermal epithelium between the left posterodorsal arm (l.pda) and the left postoral arm (l.poa), directly above the hydrocoel (hy), began to invaginate, which then became the vestibule (v) (Fig. 4A). Combination of the hydrocoel (hy) and the vestibule (v) made up the echinus rudiment (Fig. 5A). The floor of the vestibule bulged out forming five primary podia (p.p) and echinus rudiment continued to increase in size and came to dominate the left side of the larva (Fig. 5B).

The late eight-armed larva of *C. mertensii* looked ruddy and could be recognized by the naked eye. Three ciliated bands developed on the body: the preoral ciliated band (pr.ci.b) and anterior ciliated band (a.ci.b), were formed on the ventral and dorsal surfaces of the larval body, and the posterior ciliated band (p.ci.b) on the right and left sides of the posterior end of the body encircling the posterior area of the larva (Fig. 5B). On the right side of the larva were three pedicellariae (ped). The first pedicellaria was formed in the middle of the posterior part of the larva. The others were formed at the base of the right posterodorsal arm (pda) and postoral arm (poa) (see below).

In the later larval developmental stage, there were no particular differences between morphology of *C. mertensii* and *A. crassispina* except for the distribution of the pigment spots. The larval body of *C. mertensii* was vividly colored

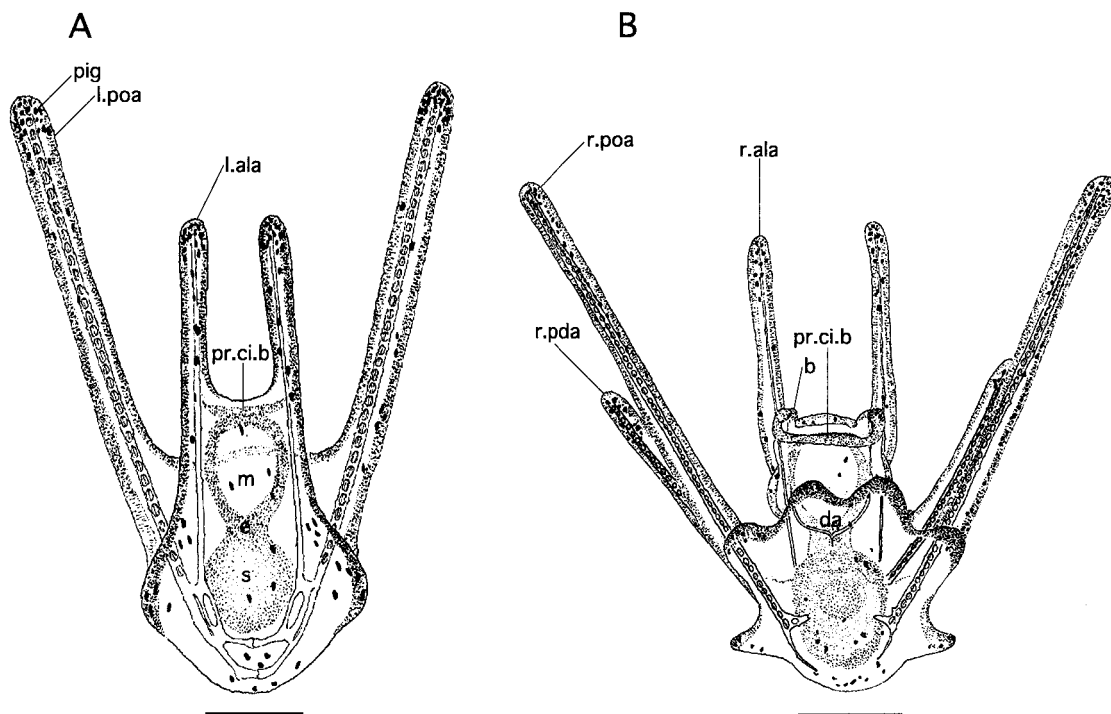


Fig. 3. Early larval stages of *Colobocentrotus mertensii*. **A.** Seven-day-old four-armed larva with well developed postoral arm and anterolateral arm. Brownish red pigment spots are observed at the tip of the larval arms. Dorsal view. **B.** Sixteen-day-old late six-armed larva with two pairs of posterodorsal arms and well-developed preoral ciliated band. Ventral view. Scale bars, 100 μ m.

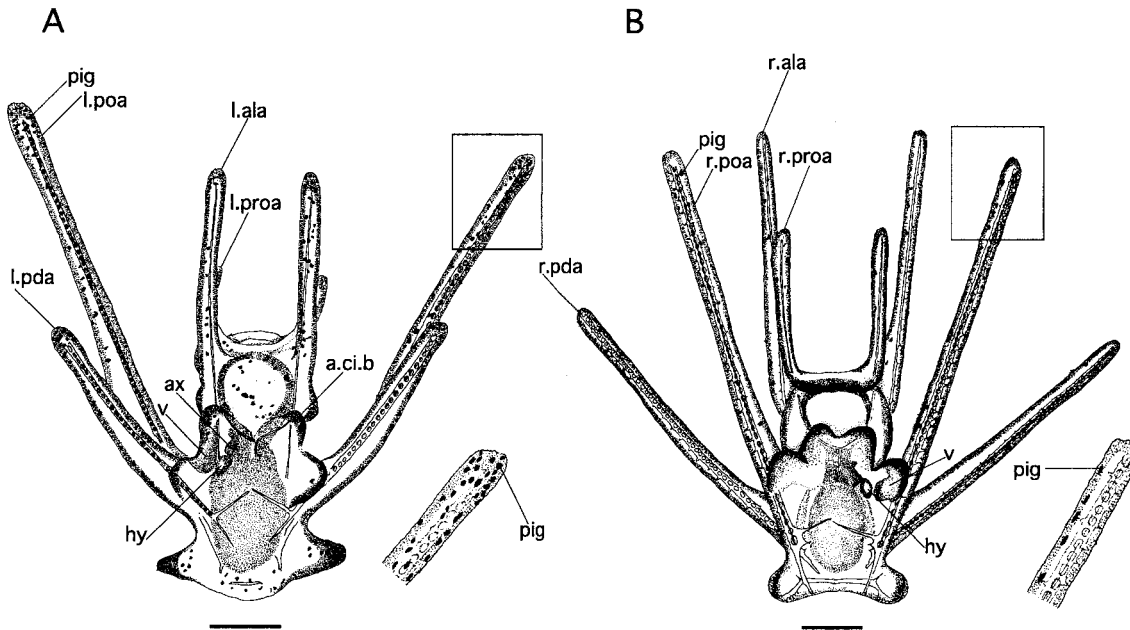


Fig. 4. Late larval stages of *Colobocentrotus mertensii* (A) and *Anthocidaris crassispina* (B). A. Twenty-three-day-old mid eight-armed larva. The hydrocoel appears as a sac on the anterior left side of the esophagus. Also on the left-hand side of the larval body is formed vestibule. Dorsal view. B. Twenty-four-day-old mid eight-armed larva. Ventral view. Tips of the arms in rectangles are shown in higher magnification to show that pigment spots are many and concentrated in *C. mertensii*, but are few and disperse in *A. crassispina*. Scale bars, 100 μ m.

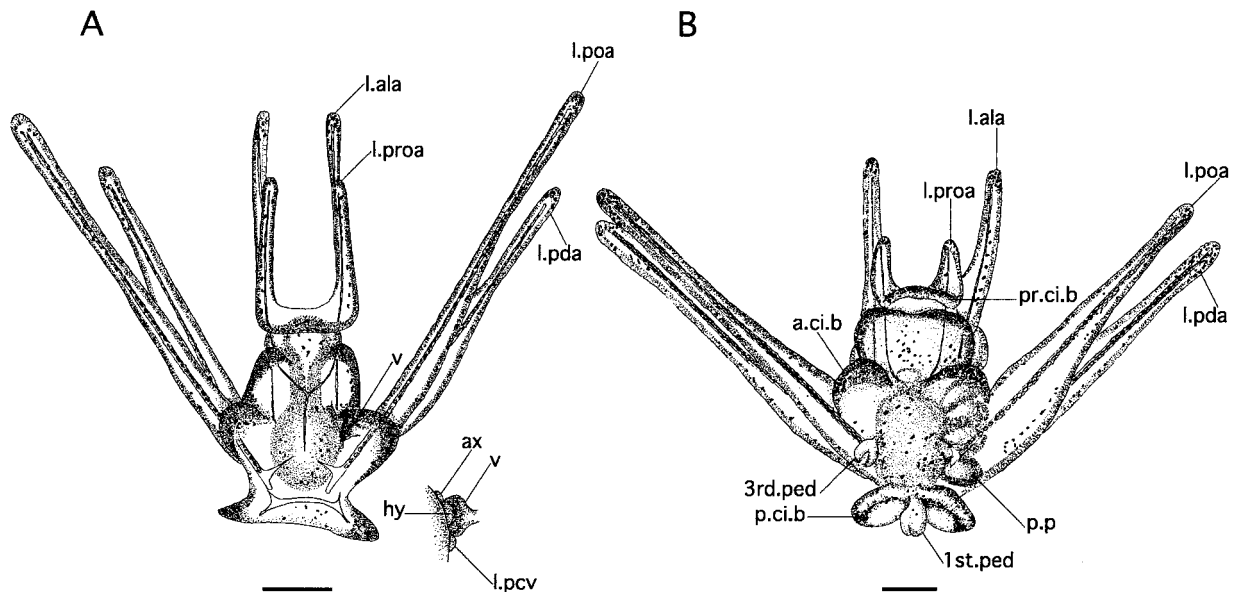


Fig. 5. Late larval stages of *Colobocentrotus mertensii*. A. Twenty-five-day-old advanced mid eight-armed larva. Echinus rudiment is developing on the left side of the body which is shown in higher magnification in an attached figure. Ventral view. B. Thirty-two-day old late eight-armed Larva. The echinus rudiment is well developed and have primary podia. Pedicellariae are present on the right side and posterior side of the body. Ventral view. Scale bars, 100 μ m.

with red pigment spots, especially at the tip of the arms (Fig. 4A). In *A. crassispina* (Fig. 4B), pigment spots were dispersed along the arms like other echinometrid larvae (Onoda, 1936).

Metamorphosis

When metamorphosis began, the anterolateral, post-

oral, posterodorsal and preoral arms were absorbed together with the skeletons and epidermis, first from left side of the larva (Fig. 6A). On the other hand, the spines covered with epithelium distended the vestibular wall and brisk primary podia (p.p) and spines (sp) protruded through the opening of the vestibular pore. Then, the primary podia and spines completely emerged to the exterior of the body. Thereafter,

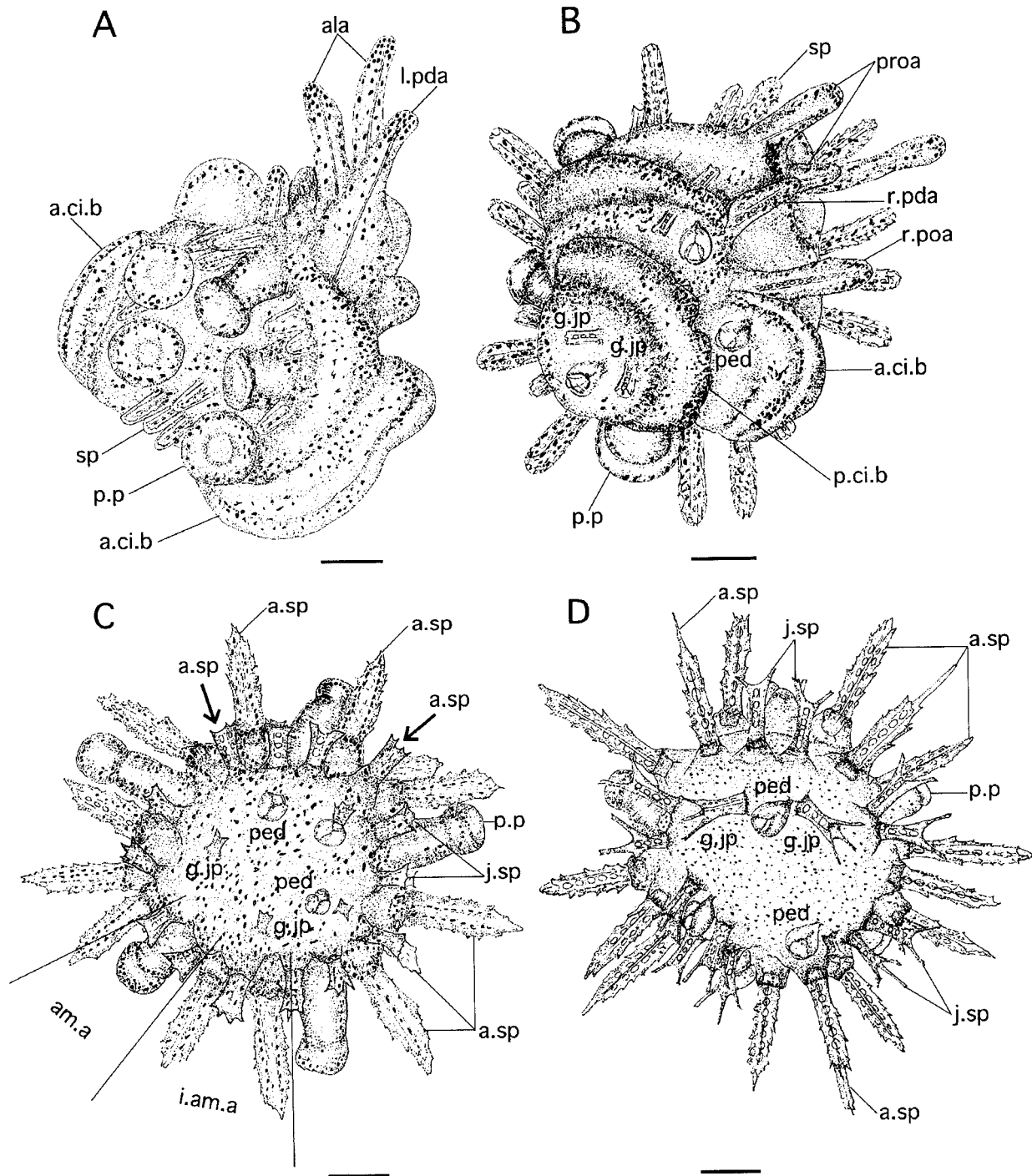


Fig. 6. Metamorphosing larva (**A**) and newly metamorphosed juveniles of *Colobocentrotus mertensii* (**B**, **C**) and *Anthocidaris crassispina* (**D**). **A.** Thirty-six-day-old metamorphosing larva with shortened larval arms. The spines and stout primary podia are protruding from the opening of the vestibule. Ventral view. **B.** Thirty-seven-day-old juvenile urchin immediately after metamorphosis. The right larval arms, ciliated bands and pedicellariae are on the aboral side, while the oral side contains adult spines and podia. Dorsal view. **C.** A juvenile two days after metamorphosis. Adult spines and juvenile spines at the circumference of the test are broad. One pair of juvenile spines alternates with each of the stout five primary podia. Arrows indicate one spine that is less developed. Dorsal view. **D.** A juvenile of *A. crassispina*, two days after metamorphosis. Adult spines and juvenile spines are more slender and their tips are sharper compared to those of *C. mertensii* (**C**). Dorsal view. Scale bars, 100 μ m.

the newly metamorphosed juvenile turned left side down so that the echinus rudiment became the lower surface and the opposite part, including the right side of postoral (r.poa) and

posterodorsal arms (r.pda), three pedicellariae (ped), and four juvenile spines (g.jp), became the upper surface of larval body (Fig. 6B). Metamorphosis took about one hour for

complete emergence of the primary podia and spines. Thus, the process of metamorphosis of *C. mertensii* was almost the same as that of *A. crassispina* (Onoda, 1931). One of the morphological features of *C. mertensii* and *A. crassispina* was found in the mode of the absorption of the arms. In these species, the epidermis and skeletons were absorbed simultaneously, whereas in some species, it has been claimed that absorption occurs only in the epidermis resulting in the naked skeletons (Emlet, 1988; Gosselin, 1998).

Juvenile structures

The form of the juvenile was penta-radial as the adult. Three pedicellariae (ped) and the juvenile spines (g.jp) were formed on the aboral surface of the juvenile, while the ciliated bands were absorbed (Fig. 6C). Juvenile morphology of *C. mertensii* began to deviate markedly from that of juveniles of other echinometrids. The juvenile of *C. mertensii* was dotted with a lot of pigment spots and became deeper in color (Fig. 1B) while that of *A. crassispina* was light brown. The aboral region was circular and five ambulacral areas (am.a) and interambulacral areas (i.am.a) were arranged alternately. In each ambulacral area, there were one central primary podium (p.p) and two broad juvenile spines (j.sp) in aboral position (Fig. 6C). Each podium was extensible and had a terminal disc of about 77 μm in diameter which was more stout and thicker than that of *A. crassispina* which had a diameter of about 62 μm with more slender disc (Fig. 6D). In the interambulacral areas were a group of four adult spines (a.sp) that began to demonstrate their adult features. The adult spines (Fig. 6C) were broader

than those of *A. crassispina* (Fig. 6D). Besides, in *C. mertensii*, one adult spine (a.sp) (shown by the arrows in (Figs. 6C and 8B) was smaller than other three spines, while in *A. crassispina*, four adult spines (a.sp) (Fig. 6D) were the same size.

Features of larval and juvenile skeletal structures

Since the formation and structure of skeletal system of larva and juvenile are important for the classification of sea urchins, we made close examination on the skeletal system of *C. mertensii* in comparison with that of *A. crassispina*.

The early larval skeleton is formed by the elongation of triradiate spicules (Okazaki, 1975). In the posterior part of the larval body develops a compound basket structure. At the later developmental stage, a new unpaired posterior transverse spicule was formed where the basket structure had disintegrated. During the mid eight-armed larva stage, posterior transverse rod (ptr) (Fig. 7A) became well developed and on each side there were two branches: one was directing upward obliquely and the other was long and directing downward. In *C. mertensii* (Fig. 7A) the lower branches were smooth, whereas in *A. crassispina* (Fig. 7B) they were furnished with a series of 5-6 thorns along their lower edge. The fenestrated postoral rod (por) and postero-dorsal rods (pdr) of *C. mertensii* had only small numbers of smooth thorns (Fig. 7A) but in *A. crassispina* they had many thorns (Fig. 7B).

The base of each postoral rod (por) expanded into a dense network (stereom) and formed a highly fenestrated plate (f.p) (Fig. 8A). A slightly smaller fenestrated plate

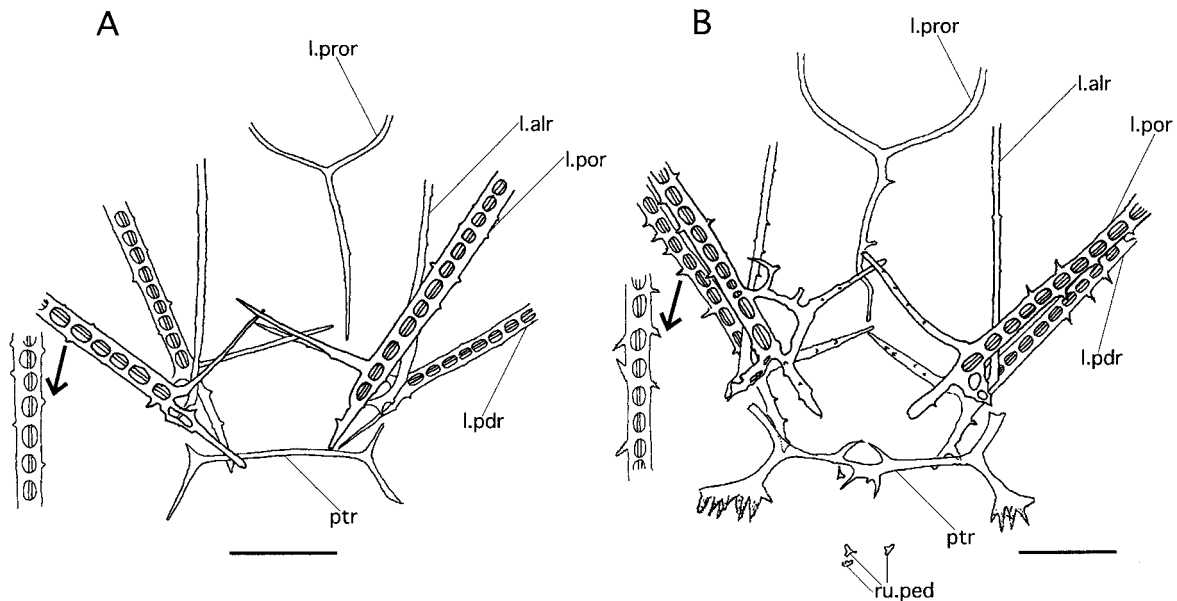


Fig. 7. Skeletal specimen of mid eight-armed larvae. **(A).** Twenty-five-day-old larva of *Colobocentrotus mertensii* **(A)** and twenty-seven-day-old larva of *Anthocidaris crassispina* **(B)**. **A** shows that two pairs of fenestrated skeletal rods and bifurcated unpaired posterior transverse rod. Ventral view. **B** shows that posterior transverse rod (ptr) with two pairs of small posterior processes and lower branches were furnished with a series of 5-6 thorns along their lower edge. Three triradiate spicules, presumptive jaws of the pedicellaria, are visible in the posterior region. Ventral view. Note that skeletal rods possess smooth thorns (arrow) in *C. mertensii*, but many sharp thorns (arrow) in *A. crassispina*. Scale bars, 100 μm .

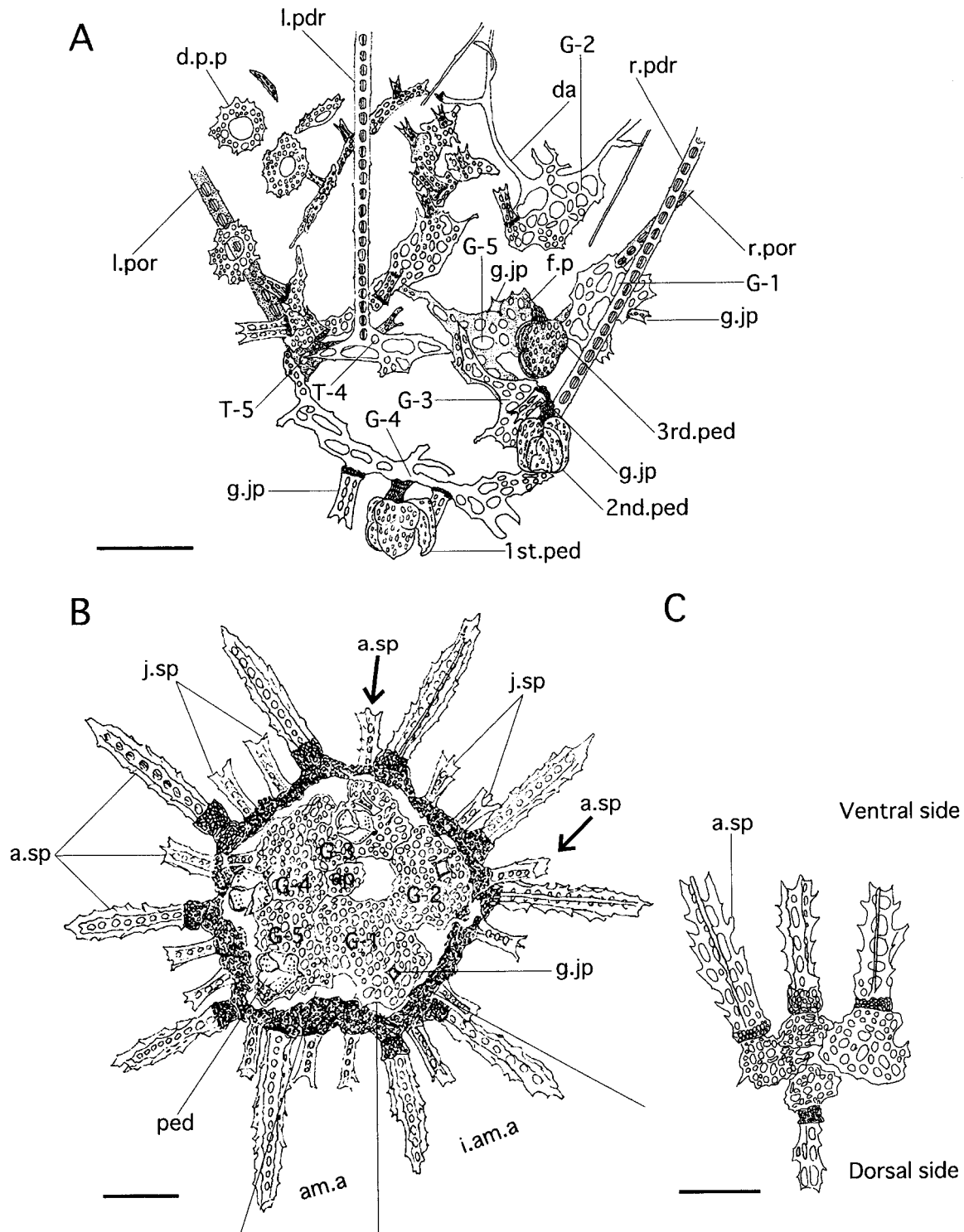


Fig. 8. Skeletal specimen of late eight-armed larva and juvenile of *Colobocentrotus mertensii*. **A.** Thirty-five-day-old larva, showing five genital plates (G-1~G-5) and two terminal plates (T-4, T-5). G-1 develops from new triradiate spicules and other plates from the larval skeletal elements. Terminal plates are formed at the base of left posterodorsal and postoral rod. Dorsal view. **B.** A juvenile two days after metamorphosis. The aboral surface of juvenile is occupied with five genital plates (G-1~G-5) and an anal plate (ap). Four broad adult spines in interambulacral area form a lozenge. Arrows indicate one of the four adult spines that is smaller in size than others. Juvenile spines are formed in ambulacral area. Dorsal View. **C.** A set of the adult spines in an interambulacral area. One adult spine located in the dorsal side is smaller in size than other three spines. Scale bars, 100 μm .

Table 2. Major differences in morphological features between *Colobocentrotus mertensii* and *Anthocidaris crassispina* in development.

Developmental stage	Morphological features	<i>C. mertensii</i>	<i>A. crassispina</i>
Eight-armed larva	Pigment spots of arms	Concentrated on the tip	Dispersed
	Thorns of skeletal rods	Smooth	Sharp
Juvenile	Color	Deep red	Light brown
	Primary podia	Stout and thick	Thin
	Juvenile spines	Broad and slightly flat tip	Slender and sharp tip
	Adult spines	Broad and slightly flat tip	Slender and sharp tip
		One adult spine is smaller than other three spines	Four adult spines are the same size

developed at the base of each posterodorsal rod (pdr). Together, the fenestrated plates originating from the postoral rods (por) and the posterodorsal rods (pdr) formed a truncated pyramidal structure. At the posterior end of the larva, first pedicellaria (1st.ped) (Figs. 5B and 8A) and a pair of juvenile spines (g.jp) (Fig. 8A) were formed in association with the posterior transverse rod. The second pedicellaria (2nd.ped) and juvenile spine (g.jp) (Fig. 8A) were formed at the base of the right posterodorsal rod (r.pdr), and the third pedicellaria (3rd.ped) (Figs. 5B and 8A) and juvenile spine (g.jp) (Fig. 8A) were formed at the base of the right postoral rod (r.por). Later on, these skeletal networks were integrated into the test of the juvenile where they formed the G-3 and G-5 genital plates (Fig. 8B). In the same way, the future G-2 genital plate developed at the basal part of the dorsal arch (da). Finally, the G-4 plate developed at the base of posterior transverse rod and the G-1 plate in the center of the right lateral field (Fig. 8A). The terminal plates (T-4, T-5) were formed at the base of the skeletal rods sustaining, respectively, the left posterodorsal (l.pdr) and postoral arms (l.por).

The other major parts of juvenile appendages, such as terminal plates (T-1, T-2, T-3), coronal plates, spines, primary podia and the five teeth of Aristotle's lantern, developed in the echinus rudiment.

During the metamorphosis, the coronal plates, ambulacral plate and interambulacral plate came to contact and formed the outline of a rigid test. In a 2-day juvenile urchin, the aboral area included five genital plates (G-1~G-5) and two types of juvenile and adult spines. These plates (G-1~G-5) were attached to one another and touched the coronal plates (Fig. 8B). These plates also delimited the future periproctal zone, where the first anal plate (ap) appeared. Four adult spines (a.sp) developed on each network plate among which one located in the dorsal side was smaller in size than the other three spines (Fig. 8B, C). These four plates formed a lozenge: one apical plate, two median plates and one basal plate formed one interambulacral area (i.am.a). Two juvenile spines (j.sp) developed on each fenestrated plate, formed an ambulacral area (am.a). These two juvenile spines (Fig. 8B) were similar to their homologous on the genital plates (g.jp) (Figs. 6B, C and 8A) regard-

ing their development and shape.

We studied the development of *C. mertensii* from early larval stages to the juvenile, comparing to those of *A. crassispina*. External features and fundamental structures of larval skeletons were similar in these two species up to the early larval stages. However, there exist some differences in later stages of development which are summarized in Table 2.

DISCUSSION

The family Echinometridae includes six species according to the classification by Shigei (1986) and commonly possesses the following morphological characters. (1) The valves of the globiferous pedicellariae have one unpaired lateral tooth. (2) The larval skeleton shows a compound basket structure. (3) The optical crystal axis of calcite of coronal plate is perpendicular to the plate of test (Shigei, 1974). However, the shape of spines demonstrates variety among them. *A. crassispina*, *Echinometra mathaei*, *Echinostrephus aciculatus* and *Echinostrephus molaris* have needle-like spines, whereas other two species, *C. mertensii* and *Heterocentrotus mammillatus*, have peculiar spines. In *H. mammillatus*, some spines on the aboral side are long, thick and very strongly developed like a pencil, while others are short and flat-topped forming mosaic over the surface (Hyman, 1955; Matsuoka, 1989). In *C. mertensii*, spines on the aboral side are short and flat-topped and look like pavement stones. Some spines around the edge of the test are broad, flat and chisel-shaped (Fig. 1A). These differences in adult forms of Echinometrid species raise the question whether they are truly in the same family.

The developmental process of larval structure of *C. mertensii* is very similar to that of *A. crassispina*. Additionally, we compared the morphological features in three species of echinometrids larvae and found that the larval structure of *C. mertensii* resembles those of other echinometrid sea urchins. Especially, it is to be noted that triradiate spicules, the first sign of larval skeleton, were formed at the mesenchyme blastula stage in *C. mertensii* (Fig. 2A) and other Echinometrid urchins including *A. crassispina*. In many other species, i.e., *Echinostrephus molaris*, *Strongylocentrotus pulcherrimus*, *Toxopneustes pileolus* and *Pseudocentro-*

tus depressus (Onoda, 1936, Fig. 2B), the triradiate spicules appeared after gastrulation. Thus, from the developmental viewpoints, larvae of Echinometridae family members share the common fundamental characteristics.

In newly metamorphosed juvenile of *C. mertensii* (Fig. 6B) and *A. crassispina*, larval arms were absorbed together with the skeletons and epidermis. On the contrary, in *Eucidaris thouarsi* (Emlet, 1988) and *Paracentrotus lividus* (Gosselin, 1998), tissue resorption is achieved by the retraction of only epidermis resulting in the naked skeleton. The naked skeletal rods will eventually be broken down. This discrepancy may be due to the species difference.

Considerable changes in the morphology of the spines of *C. mertensii* occurred in juvenile stage: they began to show peculiar features of adult spines. The adult spines (a.sp) of *C. mertensii* (Figs. 6C and 8B, C) were broader than those of *A. crassispina* which were slender with needle-like tips (Fig. 6D). These changes of spine form may reflect the habitats and foods of larvae and juveniles. However since we cannot find juveniles in the sea, the relationship between the spine form and environment must be analyzed in future. It is necessary to rear the juveniles of *C. mertensii* to adult to study the transformations of the juvenile's spines into the adult ones and to find out which structures are specific to *C. mertensii*. Although we attempted to rear the juveniles by usual method used for regular urchins, we were unable to culture them until adulthood. For that purpose it is very important to know exactly the juvenile's nutritious food.

Adult *C. mertensii* has an unusual morphology in that the spines around the periphery of the test are long but orally-aborally flattened and tube-feet are exceptionally well developed. This morphological peculiarity may reflect the behavior of this urchin. *C. mertensii* is found in wave-swept intertidal shores, where it clings to the fully exposed surface of the rocks and moves to forage at high tide. In contrast, the juvenile is similar to the other regular echinoid urchins except peculiar spines and stout primary podia, as stated above. A study on the development of *C. atratus* by Mortensen (1943) reported that there was a relationship between the reduction of spines in *C. atratus* and the hydrodynamic force of the sea waves.

Four adult spines grew from each interambulacral area (i.am.a) and one of them was smaller in size than others (Figs. 6C and 8B, C), while in *A. crassispina*, four adult spines were the same in size (Fig. 6D). In *Echinus miliaris* (Goldern, 1926), *Hemicentrotus pulcherrimus* (Noguchi, 1988), *Mespilia globulus* (Onoda, 1936), and *P. lividus* (Gosselin, 1998) four adult spines are the same size as well. Whether the small sized adult spine of *C. mertensii* will grow into the same size as other three spines or not awaits the future study.

In conclusion, this study demonstrates that the development processes of larval structure of *C. mertensii* are similar to those of members of Echinometrids including *A. crassispina*. In these urchins, the first larval spicules were formed at the mesenchyme blastula stage much earlier than in non-echinometrid species. Among echinometrids, *C.*

mertensii shows its peculiar form of spines from juvenile stage on.

ACKNOWLEDGMENTS

We are deeply grateful to Dr.Sadao Yasugi for his kind encouragement and help in preparing the manuscript. We are also thankful to Drs. Tomio Yanagisawa, Katsumi Matsuura, Isao Uemura, Makoto Kurokawa, Hidetoshi Kato and Hirobumi Suzuki. We would like to thank the staff of Tateyama Marine Laboratory, Ochanomizu Women University, for providing animals used in this study.

REFERENCES

- Emlet RB (1988) Larval form and metamorphosis of a "primitive" sea urchin, *Eucidaris thouarsi* (Echinodermata: Echinoidea Cidaroida), with implications for developmental and phylogenetic studies. *Biol Bull* 174: 4–19
- Fukushi T (1960) The external features of the development of the sea urchin, *Glyptocidaris crenularis* A. Agassiz. *Bull Mar Biol Sta Asamushi Tohoku Univ* 10: 57–63
- Gordon I (1926) The development of the calcareous test of *Echinus miliaris*. *Philos Trans Roy Soc Lond Ser B* 214: 259–312
- Gosselin P, Jangoux M (1998) From competent larva to exotrophic juvenile: a morphofunctional study of the perimetamorphic period of *Paracentrotus lividus* (Echinodermata, Echinoidea). *Zoomorphology* 118: 31–43
- Hyman LH (1955) The invertebrates: Echinodermata. In: "The Coelomate Bilateria, Vol 4" Ed by EJ Boell, McGraw-Hill Books, New York, pp 1–763
- MacBride EW (1903) The development of *Echinus esculentus* together with some points in the development of *E. miliaris* and *E. acutus*. *Philos Trans Roy Soc London* 195: 285–337
- MacBride EW (1914) The development of *Echinocardium cordatum* 1. The external features of the development. *Quart J Microsc Sci* 59: 471–486
- Matsuoka N, Suzuki H (1989) Electrophoretic study on the phylogenetic relationships among six species of sea-urchins of the family Echinometridae found in the Japanese waters. *Zool Sci* 6: 589–598
- Mortensen TH (1921) Studies on the development and larval forms of echinoderms. GEC Gad, Copenhagen
- Mortensen TH (1943) Camarodonta. In "A Monograph of the Echinoida. Vol. 3", CA Reitzel, Copenhagen, pp 277–4393
- Noguchi M (1988) External feature and body skeleton in late development of *Hemicentrotus pulcherrimus*. Ph.D Dissertation, Tokyo Metropolitan University
- Onoda K (1931) Notes on the development of *Heliocidaris crassispina* with special reference to the structure of the larval body. *Mem Coll Sci Kyoto Imp Univ (B)* 7: 103–135
- Onoda K (1936) Notes on the development of some Japanese echinoids with special reference to the structure of the larval body. *Jpn J Zool* 6: 637–654
- Okazaki K (1975) Normal development to metamorphosis. In "The Sea Urchin Embryo. Biochemistry and Morphogenesis" Ed by G Czihak, Springer-Verlag, Berlin, pp 177–232
- Shigei M (1974) Echinoids. In "Systematic Zoology Vol. 8b" Ed by T Uchida, Nakayama Book Co., Tokyo, pp 208–332 (In Japanese)
- Shigei M (1986) The Sea Urchins of Sagami Bay. Maruzen, Tokyo, pp 1–204

(Received September 3, 2003 / Accepted November 7, 2003)