CONTRIBUTIONS IN SCIENCE

A NEW BROODING BRITTLE STAR FROM CALIFORNIA (ECHINODERMATA: OPHIUROIDEA: AMPHIURIIDAE)

GORDON HENDLER AND CARLA J. BUNDRICK
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A NEW BROODING BRITTLE STAR FROM CALIFORNIA
(ECHINODERMATA: OPHIUROIDEA: AMPHIURIDAE)

GORDON HENDLER¹ AND CARLA J. BUNDRIK²

ABSTRACT. Amphiodia akosmos n. sp. is the only Amphiodia species known to bear live young, and like many brooding brittle stars, it is of diminutive size. It is characterized by irregularly arranged opaque disk scales, primary disk plates that are inconspicuous or lacking, wing-shaped middle arm spines, and a single elongate tentacle scale. There is an allometric relationship between arm length and disk diameter. Specimens of different sizes cannot be characterized by a unique ratio of arm length to disk diameter, a complication generally ignored in previous descriptions of ophiuroids. Adult individuals attain approximately 4 mm disk diameter and closely resemble the young of a larger, sympatric species, Amphiodia occidentalis. This suggests that A. akosmos is progenetic. However, various of its characteristics may be paedomorphic, perrormorphic, and nonheterochronic in origin. In contrast to female A. occidentalis, which broadcast-spawn thousands of small eggs, A. akosmos individuals hold fewer than 30 yolky eggs, which can reach 430 µm in diameter. All dissected specimens (N = 8) were female, an indication that the species could be gonochoric, but also that it might be protandric or parthenogenetic. Thus far, the species is known only from the intertidal zone within 200 km of the Monterey Peninsula, California.

INTRODUCTION

"It is a fabulous place: when the tide is in, a wave-churned basin, creamy with foam, whipped by the combers that roll in from the whistling buoy on the reef. But when the tide goes out the little water world becomes quiet and lovely. The sea is very clear and the bottom becomes fantastic with hurrying, fighting, feeding, breeding animals..." (Steinbeck, 1945)

From time to time, an unnamed species of Amphiodia has been recognized as inhabiting the central California coast. It is named and described herein, based primarily on specimens collected from the Great Tide Pool, at Point Pinos, on the tip of the Monterey Peninsula, the locality vividly depicted in the epigraph.

That naming this species has been so long delayed, despite its having been noted in two editions of Light's Manual (Weesner, 1954; Sutton, 1975), is indicative of the scant attention that has been directed toward Eastern Pacific ophiuroids. Although the first account of the fauna was published over 150 years ago (Müller and Troschel, 1843), Eastern Pacific ophiuroids have never been extensively collected, their representation in museums is limited, and the group has not been monographed. The present study is a small step toward correcting that situation.

Friedman (1948) may have been the first to note the new species. He reported two specimens, which were from Monterey Bay, of Amphiodia with "one broad tentacle scale" and a disk diameter of about 2 mm. Although he mistakenly identified them as a tropical species, his description of the animals suggests that he had found either the new species reported herein, or juvenile individuals of Amphiodia occidentalis (Lyman, 1860). Although some of Friedman's ophiuroids are deposited at the California Academy of Sciences, these specimens of Amphiodia tabogae are believed lost.

The new species was certainly known to Weesner (1954), who differentiated "Amphiodia sp." from A. occidentalis in the second edition of Light's Manual. She distinguished the new species based on its arm/disk ratio and single tentacle scales, and more definitively by its "ovoviviparous" reproduction. Unfortunately, no record exists of where along the central California coast her specimens were collected (F. Lechlieitner,née Weesner, personal communication).

The second author of the present report (under the name Cunningham, 1977) rediscovered the small, brooding Amphiodia species in the course of research on A. occidentalis. She noted additional features distinguishing the two Amphiodia species, in addition to those taxonomic characters cited by Weesner (1954). Specimens that she collected were examined by Sutton (1975) and treated as "Amphiodia sp." in the third edition of Light's Manual. However, as noted below, Sutton evidently confused juvenile A. occidentalis and adults of the new species and incorrectly concluded that the new species "May be a variety of A. occidentalis" (Sutton,

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1975:633). Until this report, nothing further was published on the species, save for references to Cunningham's observations (Austin and Hadfield, 1980; Rumrill and Pearse, 1985; Hendler, 1991).

MATERIALS AND METHODS

Standard methodology and terminology were used for the taxonomic analysis. The diameter of the disk (dd) was measured from the outer edge of the radial shields to the opposite edge of the disk. The length of the arm (AL) was measured for the longest arm of an individual, from the edge of the disk to the arm tip. Data were not recorded for specimens with broken or regenerated arms.

The AL and dd of a range of sizes of A. occidentalis from several localities were measured for comparison with individuals of the new species. The allometric equation, $Y = aX^b$ (where $Y = AL$, $X = dd$), was used to describe the relationship between arm and disk dimensions. Statistical analysis of the data was based on a Model I regression of the linearized expression, $\ln Y = \ln a + b \ln X$. Two-tailed Student's $t$ tests were used to determine the significance of the regression based on deviation of the correlation coefficient from zero, and departure from isometry based on deviation of the regression coefficient (= constant of allometry) from unity (Sokal and Rohlf, 1995).

The following abbreviations are used in this paper. CASIZ, California Academy of Sciences; LACM, Natural History Museum of Los Angeles County; USNM, National Museum of Natural History.

SYSTEMATIC ACCOUNT

Family Amphiuridae

Genus Amphiodia Verrill, 1899

Amphiodia (Amphiodia) akosmos new species

Figures 1-4


ETYMOLOGY. The specific name, transliterated from the Greek term for "unorganized," refers to the irregular distribution of dorsal disk scales and to the apparent absence of the rosette of primary disk plates that comprise an orderly system ("kosmos") in other species of Amphiodia.

MATERIAL EXAMINED. Except as noted, all specimens in the type series are preserved in ethanol and were collected from the intertidal zone in the Great Tide Pool at Point Piños, Pacific Grove, Monterey County, California, located approximately 36° 37.7'N, 121° 56.20'W.


**DESCRIPTION OF HOLOTYPE.** Disk diameter 3.3 mm; AL 16.8 mm.

**Dish.** Rounded pentagonal, lacking mid- Interradial notches, covered by irregularly arranged, imbricating scales; boundary between dorsal and ventral disk scales not sharply demarcated by closely aligned marginal scales. Primary plates inconspicuous or lacking. Scales opaque, somewhat irregularly shaped, with rounded edges; largest in dorsal interradial region, smallest in ventral interradii. Genital scales inconspicuous, tilting into bursal slit, appearing similar in size to ventral interradial scales. Bursal slit extending from near posterior margin of oral shield almost to edge of disk.
Radial Shield. Nearly semicircular, narrowing proximally, with broad, straight distal end; length equaling approximately one third of disk radius. Peripheral scale capping distal end of radial shield, subquadrate, thick, width equaling or exceeding one-half width of radial shield. Paired radial shields in contact distally; proximal ends separated by broad wedge-shaped scale, and frequently by narrow wedge-shaped scale distal.

Jaw. Narrow, with three, bluntly rounded oral papillae. Oral slits wide. Paired infradental papillae largest, blocklike, not markedly inflated, tapering distally, tilting upward toward teeth. Distal papillae erect, tabulate. Middle papilla smallest, elongate, with distal edge truncate, broader than base. Outer papilla elongate, with tapered, bluntly rounded distal edge.

Oral Shield. Rhombic, with bluntly rounded corners, width equaling or exceeding length. Proximal edges somewhat convex, longer than distal edges. Madreporite larger, rounder than oral shield; distal side expanded, convex, with single pore.

Adoral Shield. Long, narrow proximal lobe aligned with oral shield; short, distal lobe abutting first ventral arm plate. Paired shields touching within, or nearly so.

Arm. Constricted basally, gradually widening over approximately one-quarter total length, tapering to slender distal tip; broader than high in cross section; convex dorsally, flat ventrally.

Dorsal Arm Plate. Subhexagonal, twice as wide as long. Proximal edge convex; proximolateral edge meeting somewhat shorter posterolateral edge forming rounded projection; distal edge indented medially. Proximal plates slightly overlapping. Distal plates three-sided, separated by lateral arm plates, with straight obtusely angled proximal edges meeting at sharp apex, convex distal edge.

Lateral Arm Plate. Slender, with protruding articulation ridge. Dorsal side forming small, triangular wedge between dorsal arm plates, ventral side forming larger narrow wedge separating ventral arm plates. Successive plates joined by sheath of integument.

Ventral Arm Plate. Subpentagonal, width equaling or exceeding length. Proximal portion relatively thickened. Proximal edges somewhat concave, meeting at sharp obtusely angled apex; lateral edges convex, excavated to accommodate tentacle scale; distal edge longest, with medial indentation demarcating two distolateral lobes. Distal plates pentagonal, straight sided. First ventral arm plate inclining into oral slit, nearly perpendicular to ventral surface of disk; portion within oral slit, thin, narrow; outer portion C-shaped, thickened, forming flat shelf between adoral shields.

Arm Spines. Erect, three in number (except two on first joint beneath disk); length of spines approximately equal, slightly less than width of ventral arm plate. At base of arm: dorsal spine proximodistally compressed, broader than thick, with somewhat concave distal side, outer tip bluntly rounded to flared; middle spine wing-shaped in dorsal view, base triangular in cross section, with distal edge dorsoventrally compressed, keel-shaped; outer portion of spine proximodistally compressed, ovoid to rounded in cross section, tapering to bluntly rounded tip; ventral spine resembling dorsal spine though less broad, narrower at outer tip, not distally concave. Spines increasing slightly in length from dorsal to ventral.

Tentacle Scale. Single, elongate, ovoid to somewhat bilobate; proximal end frequently slightly widened, inserted into angle between lateral and ventral arm plates; adradial edge aligned with excavaed shelf on lateral edge of ventral arm plate; distal tip bluntly rounded. Length on arm joints beneath disk approximately one-half that of associated ventral arm plate.

Terminal Arm Plate. Strawberry shaped, with 0–2 sharp, minute protrusions at terminal pore.

Tube Foot. Shaft tapering, translucent; tip spheroidal, with two to four pairs of subterminal, ventrolateral, glandular bumps. Whitish in alcohol.

Color. Dorsally mostly whitish to pale beige; disk scales opaque, concealing dark brown stomach; some disk scales gray or brown, radial shields dark gray proximally, whitish distally; portions of some dorsal and lateral arm plates dark gray or black. Ventrally, some ossicles of oral frame pale gray, arm plates with whitish edges and traces of pale gray. Some proximal tube feet with pale yellowish pigmentation. All but gray and black coloration quickly lost in ethanol, most specimens fading to white.

Variation. The dd of available specimens, except embryos, ranges from 2.0–4.2 mm, AL from 13.3–23.8 mm and the AL/dd ratio from 5.0–6.9. Deviations from the holotype are manifested in the number of pores in the madreporite, which may be zero or one, the shape of the dorsal arm plates, which have a median indentation in the holotype but lack a conspicuous notch in most other specimens, and the number of wedge-shaped scales separating the radial shields. Rarely, tips of ventral arm spines are flared, and basal ventral arm plates are separated by lateral arm plates. In the smaller specimens, length and width of second ventral arm plate are equal, buccal scales (sensu Hendler, 1978)

Figure 3 Amphiodia akosmos and Amphiodia occidentalis. A–C, adult Amphiodia akosmos, 3.5 mm dd, holotype, LACM 1993-035.004; D–F, juvenile Amphiodia occidentalis, 3.4 mm dd, LACM 1993-036.007; G–I, adult A. occidentalis, 8.1 mm dd, LACM 1998-089.010. For each specimen, shown from left to right are: the arm base and adjoining portion of the disk in dorsal view, the arm base and adjoining portion of the disk in ventral view, the oral frame and adjoining portions of disk and arm. Scale a = 0.5 mm (A–F); scale b = 1.0 mm (G–I)
Table 1. Features distinguishing adult *Amphiodia akosmos* from juvenile *Amphiodia occidentalis* of similar size (2.5–3.6 mm dd).

<table>
<thead>
<tr>
<th>Species</th>
<th><em>Amphiodia akosmos</em></th>
<th><em>A. occidentalis</em></th>
</tr>
</thead>
<tbody>
<tr>
<td>Disk scolation</td>
<td>Primary plates inconspicuous or lacking, scales irregularly arranged, opaque, relatively large; scales on dorsal and ventral sides not sharply demarcated at edge of disk</td>
<td>Primary plates conspicuous, scales symmetrically arranged, thin, relatively small; scales on dorsal and ventral sides sharply demarcated at edge of disk</td>
</tr>
<tr>
<td>Peripheral scale capping distal edge of radial shield</td>
<td>Width &gt;½ width of radial shield</td>
<td>Width &lt;½ width of radial shield</td>
</tr>
<tr>
<td>Wedge-shaped scales between radial shields</td>
<td>1–2 conspicuous scales; inner scale broad, outer scale slender</td>
<td>0–1 small scales</td>
</tr>
<tr>
<td>Arm spines</td>
<td>Robust</td>
<td>Relatively slender</td>
</tr>
<tr>
<td>Middle arm spine</td>
<td>Wing-shaped in dorsal view, with basal portion dorsoventrally compressed, forming a distal keel</td>
<td>Lacking a conspicuous distal keel</td>
</tr>
<tr>
<td>Second ventral arm plate</td>
<td>Width ≥ length</td>
<td>Width ≥ length</td>
</tr>
<tr>
<td>Infradental papilla</td>
<td>Inclined toward teeth</td>
<td>Perpendicular to dental plate</td>
</tr>
<tr>
<td>Oral shield</td>
<td>Width ≥ length</td>
<td>Width &lt; length</td>
</tr>
<tr>
<td>Middle oral papilla</td>
<td>Truncate, broadest at distal tip</td>
<td>Bluntly rounded, broadest subterminally</td>
</tr>
<tr>
<td>Ventral arm plate</td>
<td>Distal edge bilobate, with medial indentation</td>
<td>Distal edge straight or slightly convex</td>
</tr>
<tr>
<td>Number of tentacle scales</td>
<td>1</td>
<td>0–2</td>
</tr>
<tr>
<td>Tentacle scale</td>
<td>Length &gt;½ length of ventral arm plate</td>
<td>Length ≤½ length of ventral arm plate</td>
</tr>
<tr>
<td>Genital scale</td>
<td>Inconspicuous</td>
<td>Conspicuous</td>
</tr>
</tbody>
</table>

are present that are later resorbed, and the broad abradial edge of the radial shield is bare of scales. In some specimens the dorsal disk scales are pale brown and the outer tips of the arm spines are pale brown. Features of a newly emerged juvenile are considerably different from those of adults. It had five arm joints, which lacked tentacle scales, and its arms (0.46 mm) were shorter than the dd (0.67 mm).

**COMPARISONS.** A.M. Clark (1970) noted that only four of the nominal species in the subgenus *Amphiodia*, from east or west America, have three approximately equal oral papillae on each half of the jaw, as opposed to an enlarged operculate outer oral papilla. All except *Amphiodia pulchella* (Lyman, 1869) and *A. akosmos* have two tentacle scales. *Amphiodia pulchella* is readily distinguished from *A. akosmos* by its very finely scaled disk and slender radial shields.

Of the remaining eight species that are considered valid members of *Amphiodia* by A.M. Clark (1970), *Amphiodia euryaspis* H.L. Clark, 1911, must be referred to *Amphioptus* (see Sutton, 1976). The three congers that, like *A. akosmos*, have equal oral papillae and a single tentacle scale are differentiated from it as follows. *Amphiodia acutispina* Koehler, 1914, has a thorny projection from the outer edge of the radial shield, and both *Amphiodia minuta* H.L. Clark, 1939, and *Amphiodia obtecta* Mortensen, 1940, have slender, acutely pointed arm spines. Two additional *Amphiodia* species were described subsequent to A.M. Clark’s (1970) monograph. *Amphiodia guillermosoberoni* Caso, 1979, is distinguished from *A. akosmos* by its paired tentacle scales and elongate, narrow radial shields. *Amphiodia picardi* Cherbonnier and Guille, 1978, has an *Amphioptus*-type oral formula and is probably an *Ophionephtys* species.

**Distinctions Between Adults and Juveniles of *A. akosmos* and *A. occidentalis*.** Despite their similarities, *A. akosmos* is readily distinguished from the much larger adult individuals of co-occurring *A. occidentalis* (Figs. 1, 3). *Amphiodia occidentalis* reaches 12 mm dd with AL/dd ratios of up to 18, and adult individuals are more richly pigmented than *A. akosmos*, with considerable gray and brown dorsally and with dark gray on the ventral surface of the arms (e.g., see Austin and Hadfield, 1980: fig. 10.4). Furthermore, adult *A. occidentalis* consistently have two tentacle scales, ventral arm plates that are distinctly wider than long, dorsal arm plates that are 2.5 to 3 times wider than long, and radial shields that are narrow, not semicircular in shape. There are also marked differences in the reproductive systems of the two species, as noted below.

Juvenile *A. occidentalis* that are equivalent in size to *A. akosmos* are only distinguishable upon microscopic examination (Table 1; Figs. 2, 3). Among their most noticeable differences is that the disk of *A. akosmos* lacks noticeable primary plates and has somewhat larger scales than juvenile *A. occidentalis*.
Figure 4 Amphiodia akosmos and Amphiodia occidentalis. Allometric relationships between the dd and AL (top left) and the AL/dd ratio (top right) for the full size range of the specimens examined. Scatter plots of the relationship between AL and dd (bottom left) and the AL/dd ratio (bottom right) for the specimens in the size range of A. akosmos (0.5-4.5 mm dd)

(Fig. 2). Nielsen's (1932) statement that A. occidentalis characteristically lacks primary plates is incorrect.

Although individuals of A. occidentalis with regenerating disks do not have primary plates, they would not be confused with A. akosmos since, as noted during this study, it was only large adult individuals of A. occidentalis that had regenerating disks. Amphiodia akosmos has considerably more robust arm spines than juveniles of A. occidentalis, and its middle spine is noticeably wing-shaped (Fig. 2). In addition, juvenile A. occidentalis has zero to two tentacle scales, but A. akosmos has a single, markedly elongated tentacle scale and relatively more prominent wedge-shaped scales separating the radial shields.

The possession of a single tentacle scale was previously suggested to distinguish the two species (Weesner, 1954; Sutton, 1975). However, we found that the number of tentacle scales of A. occidentalis shifted from zero to one to two in individuals of increasing body size. Those with no tentacle scales were 0.54 to 3.72 mm dd (1.75 ± 0.73 [± SD], N = 71), those with one tentacle scale were 2.01 to 3.72 mm dd (2.90 ± 0.41, N = 31), and those with two tentacle scales were 2.93 to 11.97 mm dd (5.51 ± 1.73, N = 162). Thus, individuals of A. akosmos with one tentacle scale spanned the size range of A. occidentalis with zero to two tentacle scales. Amphiodia akosmos lacked tentacle scales early in development, as noted for a 0.67-mm dd juvenile, but it appears that Sutton (1975) was mistaken in stating that it can have two tentacle scales. He likely confused A. akosmos with juvenile A. occidentalis.

Allometric Relationship Between Arm Length and Disk Diameter. The sizes of A. akosmos and A. occidentalis overlap (Figs. 1, 4), although individuals of A. occidentalis can attain a much greater size than A. akosmos, up to 12 mm dd and 164 mm AL. Although it had been suggested that the two species are distinguishable on the basis of their size and AL/dd ratio (Weesner, 1954; Sutton, 1975), that is not invariably the case. The AL/dd

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Table 2. Relationship between the AL and the AL/dd ratio and dd (body size) of Amphiodia akosmos and Amphiodia occidentalis. The allometric equation is shown as a power function and in a linearized form.

<table>
<thead>
<tr>
<th>Species</th>
<th>AL = ax^b</th>
<th>ln Y = ln a + b ln X</th>
<th>N</th>
<th>r^2</th>
<th>t(r = 0)</th>
<th>t(b = 1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. occidentalis</td>
<td>2.909(dd)^1.756</td>
<td>ln AL = 1.068 + 1.756(ln dd)</td>
<td>264</td>
<td>0.953</td>
<td>72.879*</td>
<td>31.500*</td>
</tr>
<tr>
<td>A. akosmos</td>
<td>1.607(dd)^2.177</td>
<td>ln AL = 0.475 + 2.177(ln dd)</td>
<td>21</td>
<td>0.917</td>
<td>14.495*</td>
<td>7.846*</td>
</tr>
<tr>
<td>A. akosmos</td>
<td>1.607(dd)^2.177</td>
<td>ln AL = 0.475 + 1.177(ln dd)</td>
<td>21</td>
<td>0.764</td>
<td>7.842*</td>
<td>1.180 ns</td>
</tr>
</tbody>
</table>

Abbreviations: b, the constant of allometry; t(r = 0), the calculated value of t in the test for significance of the regression, where H_0: correlation coefficient = 0 vs. H_1: correlation coefficient ≠ 0; t(b = 1), the calculated value of t in the test for significant deviations from isometry, where H_0: constant of allometry = 1 (i.e. isometry) vs. H_1: constant of allometry ≠ 1 (i.e., allometry); * P ≤ 0.05; ns, not significant.

The ratio of A. akosmos examined ranged from 0.69 to 6.89, and the ratio for A. occidentalis ranged from 0.74 to 15.79.

For A. occidentalis and A. akosmos, there was a statistically significant correlation between both AL and AL/dd ratio and body size (dd) (Table 2; Fig. 4). For both species, the coefficient of determination (r^2) for the nonlinear allometric regressions (Y = ax^b) exceeded that for linear regressions of the data (Y = a + bX). The relationships (except AL/dd vs. dd for A. akosmos) were allometric since the hypothesis that b = 1 (b = the allometric constant) was rejected (Table 2). For both species, a single ratio of AL/dd is insufficient to characterize either species.

There has been only one comparable examination of the ophiuroid arm allometry, and the results also indicated a positive allometric relationship (López Vélez, 1956). However, the same situation likely prevails in other long-armed ophiuroid species, casting doubt on assumptions in the literature that the ratio between AL and dd is linear and independent of body size.

The AL and AL/dd values for many A. occidentalis exceeded those for A. akosmos of equivalent disk size (Fig. 4). However, the difference in the values could not be reliably used to distinguish the species because the arm lengths, and AL/dd ratios, of A. akosmos and A. occidentalis of comparable body size (dd) were overlapping. Analysis of covariance tests of the significance of differences between the regression coefficients of the two species were not warranted, because of departures from normality and heteroscedasticity, even of log-transformed data.

In small individuals, values for AL and AL/dd were depressed compared to those for adults, because the production of arm joints was not initiated until after the disk formed. The smallest A. akosmos examined was 0.67 mm dd with AL 0.42 mm. The smallest A. occidentalis was 0.54 mm dd with AL 0.4 mm, but another that was 0.67 mm dd already had 1.6-mm arms.

**Biology.** Distribution, Habitat, and Associations. Amphiodia akosmos has been found at the Great Tide Pool, at Pt. Piños, for the past 25 years and possibly longer. The population at that locality is protected by an offshore reef at low tides, but subject to considerable wave surge at high tides. The species also has been found nearby at Hopkins Marine Station, where specimens were last collected in 1921; more recently at Waddell Creek, Santa Cruz County, a more southerly site; and offshore in an intertidal cove at Southeastern Farallon Island. It would not be surprising to find A. akosmos at other intertidal localities in and beyond the Monterey region if the California and Oregon coastlines were carefully surveyed.

At Pt. Piños, the type locality, A. akosmos occurred in low intertidal pools that were exposed at −0.4 ft. (−0.1 m) mean lower low water, and surrounded and protected by algae-draped, granitic boulders. Individuals were found in the shallow layer of sandy sediment beneath and around cobbles and often occurred under the same rocks as A. occidentalis and Amphipholis squamata (Delle Chiaje, 1828). As an indication of their relative numbers, more than 135 A. occidentalis were collected on 23–24 July 1993 in contrast to only seven A. akosmos. Occasionally, individuals of Ophiofroctus esmarki Lyman, 1874, were also found beneath the cobbles, and Ophiotrichia spiculata Le Conte, 1851, was associated with algae at the type locality.

In 1993, unidentified stalked protozoans were found on A. occidentalis in the Great Tide Pool. Some of the preserved specimens of A. akosmos collected there at the same time had similar organisms attached to the oral frame and the basal arm spines and tube feet.

**Reproduction.** Amphiodia akosmos broods and may be reproductive for much of the year. Brooding individuals were collected from the Great Tide Pool in May 1975 and February 1976 (Cunningham, 1977). Additional brooding individuals were found in October 1999 and February 2000 (this report). Records of the occurrence and developmental stage of the brooded embryos reinforce the notion that individuals in the Great Tide Pool population do not spawn in synchrony. They also indicate that spawning individuals shed their ripe oocytes at one
time, since the embryos in all bursae of brooding females are at the same stage of development.

One specimen of 3.5 mm dd collected in May 1975 was brooding nine embryos, and another specimen collected in October 1999 of 3.0 mm dd contained only one embryo with one to two arm joints. One individual collected in February 2000 was brooding 10 embryos with only terminal arm plates; another was brooding 11 embryos with four to six arm joints (Fig. 5). Both individuals had one to five embryos in each interradius; only one interradius was barren. There were larger oocytes in the ovaries of the individual with larger embryos, indicating that vitellogenesis occurs during brooding.

A single juvenile, with five arm joints, emerged from one brooding female collected in February 2000. It was capable of locomotion using its tube feet and was negatively phototactic (Bundrick, previously unpublished). However, its emergence was probably premature, since the juvenile's disk was still pink and inflated with stored yolk. The brooded embryos in the same female had their arms reflected dorsally, such that their tube feet and mouth were in contact with the bursal wall in a position that may facilitate the uptake of nutrient (Walker and Lesser, 1989; Byrne, 1994; Hendler and Tran, 2001).

The three individuals collected in October, two in November, and one in December 1999, and two more in February 2000 were dissected; all were female. Thus, the species could be gonochoric, but protandry and parthenogenetic reproduction cannot yet be ruled out without further study. The females had a total of 10 gonads, two within each ventral interradius; additional gonadal tubules were not observed dorsal to the arms (Fig. 5). Each ovary contained approximately one to four oocytes, and individuals appeared to have 30 or fewer pinkish-orange colored oocytes. Most of the oocytes were smaller and contained less yolk than the largest, which was 430 μm in diameter. One A. akosmos embryo with a single arm joint was found to have a 400 μm dd. It is likely that it developed from an oocyte of approximately that diameter, since early developmental stages of brooding ophiuroids are often similar in size to the oocytes from which they arise (Hendler, previously unpublished). The low fecundity and the presence of large yolky eggs noted in A. akosmos are not unusual traits among brooding ophiuroids (Hendler, 1991).
In contrast to A. akosmos, the larger, but morphologically similar, species A. occidentalis is a gonochoric broadcast spawner that releases gametes between late May and June in Monterey Bay (Rumrill and Pearse, 1985). Adult individuals were reported to have up to 67,965 yellow-green eggs with a maximum diameter of 90–106 μm, and pelagic larvae (Rumrill and Pearse, 1985; Strathmann and Rumrill, 1987). However, an unpublished study of the species carried out at Friday Harbor, Washington indicated the eggs are orange-pink in color and 190 μm in diameter and that development is benthic, rapid, and direct, producing crawling embryos that hatch from a thick, sticky fertilization envelope (Emlet, personal communication). Amphiodia occidentalis was collected from the Great Tide Pool, California, and Pt. Arago, Oregon, in June 1994. The females examined were undergoing oogenesis and had oocytes of mixed sizes. The largest ripe oocytes were pink and approximately 160–192 μm in diameter, consistent with Emlet's observations (Hendler, previously unpublished).

In the previous literature, only one nominal species of Amphiodia, A. affinis (Studer, 1885), was reported to brood embryos (Mortensen, 1936; Hendler, 1991). Its generic placement is unclear, but it is definitely not an Amphiodia (Clark, 1970; Thomas, 1975). Thus, A. akosmos presently is the only Amphiodia species known to bear live young. It would be unusual among brooding ophiuroids in having separate sexes, if it is indeed gonochoric. Among the approximately 70 known species of brooding ophiuroids only 13 species are gonochoric, but 40 species are hermaphroditic (Hendler, 1991; Hendler and Tran, 2001). The strong association between brooding and hermaphroditism among the ophiuroids remains an enigma (Hendler, 1975, 1991). The brooding mode of reproduction in this and other diminutive ophiuroids may, as previously suggested (Hendler, 1991), offset disadvantages inherent in their small size.

Behavior and Morphology. Adult individuals propel themselves by simultaneously shifting the position of their arms and walking on their tube feet. Individuals quickly retract their arms when disturbed. As it is drawn toward the disk, each arm kinks into sinuousoidal waves, which compress into figure eight-shaped loops. Loops at the base of the arm are largest, and those of adjacent arms interdigitate. The five retracted arms form a tightly coiled mass on the dorsal surface of the disk, giving the animal a spherical shape. The behavior is rapid when an individual is wafted into the water column and results in its quickly sinking to the bottom like a pebble. A similar behavior is shown by A. occidentalis.

The great morphological similarity of A. akosmos and A. occidentalis, the direct development of their embryos, and their similar behavior and overlapping ranges, suggest that they are closely related. Certain features of A. akosmos give it a juvenile appearance: its relatively short arms, small number of disk scales, single tentacle scale, narrow arm plates, and broad radial shields. This suggests that it is paedomorphic (as defined by Gould, 1977). However, other traits of A. akosmos give it the appearance of a small ophiuroid species with some adult features, and suggest that it is peramorphic. It develops large tentacle scales and conspicuous wedge-shaped scales separating the radial shields at a smaller body size than many individuals of A. occidentalis. Moreover, its relatively broad second ventral arm plate is more like those of adult than of juvenile A. occidentalis. However, its peculiarly elongated tentacle scale, robust arm spines, and relatively large, opaque, irregularly arranged disk scales differ from the features of both juvenile and adult A. occidentalis and may represent adult features or traits that arose de novo.

It is tempting to view A. akosmos as progenetic—exhibiting precocious sexual maturation and retaining the juvenile characters of its ancestral species. The same mechanism has been proposed as an agent in the evolution of other species of Ophiuroidea (e.g. Matsumoto, 1917; Hendler, 1979; Vadon, 1990). However, assessments of heterochrony require a phylogenetic framework, so that ontogenetic trajectories between ancestral and derived species can be compared. Unfortunately, the necessary framework is lacking in the putative cases of ophiuroid paedomorphosis. Furthermore, as Raff (1996) has emphasized, characteristics that are presumed to be evidence of heterochrony can be the end result of other mechanisms, which are poorly understood. Thus, a better understanding of ontogenetic processes and of phylogeny in Amphiodia species are both necessary for a satisfactory appreciation of the evolution in the genus.

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LITERATURE CITED


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