Morphology, anatomy and histology of *Doto uva* Marcus, 1955 (Opistho-branchia: Nudibranchia) from the Chilean coast

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Key words: Opisthobranchia, Nudibranchia, *Doto*, Histology, Morphology, Chilean coast

Abstract

*Doto uva* Marcus, 1955 is a nudibranch species recorded from the Brazilian and Chilean coast. In spite of its wide distribution, *D. uva* has been described only superficially, mainly as to the pattern of its coloration, external morphology, radular teeth and reproductive system. Here we substantially extend this description, paying special attention to the morphology, anatomy and histology of the digestive and reproductive system. Furthermore, new data on the morphology of the central nervous system and its cell types are given, and finally, the egg mass is described. On the basis of these data, we consider the *Doto* specimens described from Chile conspecific with those from Brazil.

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Introduction

The genus *Doto* Oken, 1815 is one of the less known nudibranch genera. According to Odhner (1936), Marcus (1961), Schmekel and Portmann (1982), McDonald (1983) and Thompson and Brown (1984), the main characters to diagnose the genus *Doto* are as follows: body limaciform and highly arched; foot narrow, linear, rounded anteriorly, tapering posteriorly to a short, blunt tail; anterior foot corners rounded and not protruding; cerata without cnidosac, bulbous, tuberculate, rather club-shaped, and arranged in dorso-lateral rows on each side, the large anterior ceratal pairs bearing a plume-like “gill” on the inner surfaces; round frontal veil instead of oral tentacles; rhinophores smooth and bluntly tapered, retracted into calyciform sheaths; anus and nephroproct in latero-dorsal position; small jaws with smooth masticatory border; radula uniseriate, with numerous, somewhat asymmetrical teeth with few denticles; penis unarmed.

At the Pacific coasts ca. 11 species of the genus *Doto* have been described, distributed over North America, South America and Japan (O’Donoghue, 1921; Baba, 1938, 1949, 1955, 1971; Marcus, 1955, 1957, 1959, 1961; MacFarland, 1966; Marcus and Marcus, 1967; Gosliner and Williams, 1970; Farmer, 1980; McDonald, 1983; Behrens, 1991; Schrödl, 1996, 2003). The characters to distinguish these species have been mainly morphological, whereas Lemche (1976) took a wider approach including not only body colour pattern but also food preference and shape of spawn. Nevertheless, descriptions of the various *Doto* species are rather fragmentary and sometimes inconsistent (Baba, 1955, 1971), indicating the poor knowledge about this genus.

The present study concerns a detailed description of *D. uva* Marcus, 1955. Originally, Marcus (1955) described and Marcus (1957) redescribed this species...
from the coast of Sao Paulo, Brazil, and later Marcus (1959) recorded it at the southern Chilean coast, in Ancud, Chiloe Island. More recently, Schrödl (1996, 2003) recorded D. uva in Coliumo (central Chilean coast) and Schrödl et al. (2005) from the southern Chilean Comau Fjord, whereas Fischer and Cervera (2005) considerably extended its known geographical distribution range, to Tongoy, Coquimbo and Tocopilla (northern Chilean coast). Although relatively widely distributed and frequently found, D. uva has been described only superficially, and the data available mainly concern its general coloration, external morphology, and the gross anatomy of the radula and of the reproductive system. Here we present a detailed study on the morphology, anatomy and histology of D. uva from the Chilean coast, with special attention to the reproductive, digestive and central nervous system (CNS), contributing in this way to the knowledge of the genus Doto in general and of Doto uva in particular.

Materials and methods

The following specimens of Doto uva were studied. (1) 25 specimens, deposited as number MZUSP-25275 in the Museo Zoologico Universidad de Sao Paulo Brazil, collected at Isla San Sebastiao, Sao Paulo, Brazil (Leg. L. Formelis); (2) 1 specimen, deposited as number SMNH-1516 in the Zoological Museum in Stockholm (Sweden), collected during the Lund Expedition 1948-1949, station number M17, on December 14, 1948 at Golfo de Ancud, Canal Calbuco, Chile (41°46′30″S, 73°06′45″W), preserved length 4.5 mm (Table 1); (3) 1 specimen, deposited as number SSUC-3038 in the ‘Colección de Flora y Fauna Profesor Patricio Sanchez Reyes, Pontificia Universidad Católica de Chile’, collected at Punta Blanca, Tocopilla, Chile (22°04′S; 70°12′W), preserved length 5 mm; (4) 6 specimens, collected on March 5, 1992, in Bay La Herradura, Coquimbo, Chile (29°58′S; 71°22′W) at a depth of 5 metres, preserved lengths 3.0-5.0 mm, and (5) 3 specimens, collected on January 4, 1995 in Bay Tongoy, Coquimbo, Chile (30°15′S; 71°30′W), at a depth of 3.5 metres, preserved lengths of 4.7-6.0 mm (Leg. M.A. Fischer).

Specimens from (1) to (3) were only observed as to their external morphology. Specimens from (4) and (5) were observed alive for their morphology, and subsequently refrigerated at 4°C for 6 hrs, and fixed in 4% formaldehyde in seawater, for 20 days, and then stored in 70% ethanol. Four specimens from the localities La Herradura and Tongoy were dissected to study their anatomy. From two of these specimens the buccal mass was dissected and subsequently immersed into 10% NaOH until the tissue surrounding the jaws and radula had been dissolved, and then the material was observed in toto under the light microscope. One complete specimen of Tongoy was dehydrated in a graded ethanol series, embedded in hydroxy-ethylmethacrylate and cut into 2.5 μm serial sections on an Leica Reichert Autocut microtome (Leica Microsystems, Wetzlar, Germany) for histological observation. Sections were stained with toluidine-blue to identify structures on the basis of metachromasia.

Results

Systematics

The systematic position of D. uva is as follows.

CLADOBANCHIA Willan and Morton, 1984
DENDRONOTOIDEA Sars, 1878
Dotidae Gray, 1853
Doto Oken, 1815 (type species: Doris coronata Gmelin, 1791) I.C.Z.N opinion Nº 697
Doto uva Marcus, 1955

External morphology

The body of D. uva is limaciform and arched, being highest in its mediodorsal area (Figs. 1A, B, 2A, B). The head shows a pair of smooth rhinophores with blunt apices, surrounded by rhinophoral sheaths that reach a third of the total length of the rhinophores in living specimens (Figs. 1A, B, 2A-B). A veil without projections is present in front of the rhinophores. The sheaths are frontally tapered and protrude more or less prominently depending on the specimen (Fig. 1A, C, D-F; 2C-E).

On each side of the body, up to 7 cerata occur (Table 1; Figs. 1A, F, 2A, B). Each ceras is connected to the body by a short peduncle (Fig. 2F), has 4-6 rings of 4-6 semiglobular tubercles and one tubercle at the top (Fig. 2F, G). Cerata are club-shaped
with tubercles that decrease in size towards the peduncle. At the inner side of a ceras a pseudobranch is situated with 3-4 ramifications of varying lengths (Fig. 2F, G). Cerata can be autotomized from the peduncle (Fig. 1B, C) in response to a threatening challenge. In some specimens, small, regenerating cerata were observed.

The genital orifice is located at the right side of
the body, below the first ceras, and protrudes as a cylindrical genital papilla (Figs. 1C, F, 2B). At this side, between the first and second cerata, the anus and nephroproct form together a prominent papilla (Fig. 2A, B). The voluminous pericardium (Fig. 2A, B) starts just behind the first ceras. The tail is short and undulated. The foot is narrow and extends slightly from under the mantle (Fig. 2H).

**Coloration**

The coloration patterns of living and preserved animals are nearly the same. The body is transparent yellow to creamish white, with dark-brown dots dispersed over the dorsum, the lateral flanks, the cephalic rhinophoral sheaths, the cerata and the anal papilla (Figs. 1, 2). The rhinophores are covered with small white dots from halfway till the apex. In addition, they show small white dots at the sheath borders.

The digestive gland is clearly visible through the transparent body wall, as a thick, darkish brown band (Figs. 1B). In specimens from La Herradura the digestive gland is nearly black but in specimens from Tongoy it is grayish. The gland starts behind the rhinophores and ends in front of the tail. The tubercles of the cerata are covered with dark-brown dots but have a black-pigmented subapical ring and a white, transparent apex. Pseudobranchia are rather dark at the rachis and light at the ramifications (Fig. 2F). The black pigment of the eyes is visible at the basis of the transparent rhinophores.

**Digestive and excretory system**

**Anatomy** (Fig. 3). The masticatory apparatus was studied in detail in two specimens (preserved length: 4 mm). The very thin jaws measure 1.7 mm in length and have smooth masticatory processes (Fig. 3A). The uniseriate radula is small, about 0.75 mm in length, and has a teeth formula 61-63 × (0.0.1.0.0). The asymmetrically arched teeth are horseshoe-shaped, with up to 5 lateral denticles and one central cusp. The number of denticles, however, varies within a given radula, and some teeth may show as few as 2 denticles. The largest tooth measured 14 μm and the smallest 11 μm (Fig. 3B, C).

**Histology.** The salivary gland consists of two main parts. The distal part, located just in front of the buccal bulb, is formed by numerous glandular cells that are violet-stained and have a small, round and deep-blue nucleus (T1). It opens into the oesophagus via a thin duct. The proximal part has a globular appearance and reveals three different cell types as judged from their size, shape and staining (Fig. 4A). The first type (T1) is similar to that of the distal part of the gland. The second type consists of medium-sized cells with a small, blue nucleus and a deep-blue nucleolus (T2), and the third type is formed by very large cells that reveal a granulated cytoplasm and a large, central, blue nucleus with a deep-blue nucleolus (T3). Finally, around the globular part, just outside the gland, we counted up to twelve giant cells (mean diameter ca. 150 μm). They are blue, with a large nucleus and several nucleoli (Fig. 4A, B).

Generally, the stomach epithelium is only slightly folded but in the direction of the intestine folds become deeper. Inside the cerata the digestive gland is highly folded. In the epidermis of the cerata intensely stained glandular cells occur (Fig. 4C). The kidney has a thin and highly vacuolated epithelium (Fig. 4E) and is connected to the pericardium by a piriform syrinx (Fig. 4D). The anus is located close to the thin nephroproct, and shows a folded and strongly ciliated epithelium (Fig. 4E).

**Pedal gland**

**Histology.** The pedal gland (Fig. 4F) lies concentrated in the anterior periphery of the foot, in the lips and around the oral opening. It is constituted of numerous semispherical follicles with deep-rose-staining cells that have a strongly granulated cytoplasm and a dark-blue, centrally located nucleus.

**Reproductive system**

**Anatomy** (Fig. 5). The anatomy of the reproductive system was reconstructed using serial sections. The ovotestis appears like a granulated, compact mass, lying over the kidney and occupying nearly one-third of the visceral cavity. It is connected with the ampulla by a short hermaphroditic duct. The ampulla is thick, refringent and globular-shaped. The short, postampullar spermoviduct splits into a male and a female part. Immediately after the bifurcation of the spermoviduct the male duct forms a sausage-shaped prostate gland and then runs into the muscular penial
bulb with its muscular, unarmed penis. The female part starts with a valve situated at the beginning of the oviduct that contacts the voluminous, piriform seminal receptacle and also the mucous gland. The very long vagina forms a U-shaped loop close to the upper part of the penial bulb. The vaginal duct becomes bulbous near its connection with the seminal receptacle, which is controlled by a small sphincter.

The female gland consists of three parts: the large mucous gland, the medium-sized capsule gland and the small membrane gland.

**Histology** (Figs 6, 7). The ovotestis consists of numerous acini. Generally, 5-6 globular, female acini surround one large, globular, male acinus. The hermaphroditic duct and the ampulla are lined by a thin and strongly ciliated epithelium. The ampulla
Fig. 3. A: Right jaw, dorsal view. B: Rachidian tooth. C: Radular teeth, lateral view. Scale bars: A: 1 mm, B,C: 10 μm.
lumen is filled with spermatocytes (Fig. 6A). The prostate gland consists of two parts. The distal part has ciliated cells of a globular shape and varying height (Fig. 6B). Their nucleus is violet and located in basal position whereas the curved cell apex faces the lumen. The proximal part of the prostate shows highly columnar cells filled with granules, and a darkish violet nucleus in basal position (Fig. 6C). Together with these cells, small, triangular, ciliated epithelial cells are present. The ciliated epithelium of the vas deferens is surrounded by muscular layers. The muscular penial bulb is very well developed and the penis is smooth (Fig. 6D).

The ciliated vaginal epithelium consists of cubical and elongate cells with a central nucleus and a vacuolated cytoplasm (Fig. 7A). The seminal receptacle shows six lobes, each of which is studded with allosperm, with the sperm heads oriented towards the epithelial wall and the tails towards the lumen (Fig. 7A, B). A long, ciliated duct emerges from the receptacle, two small valves separate the oviduct from the spermoviduct (Fig. 7C). The capsule gland (Fig. 7D) has columnar secretory cells with a basal nucleus and a granulated cytoplasm with deep-blue secretory droplets. In between these cells, small, ciliated supporting cells occur that are very narrow at the base but taper towards the apex. The membrane gland (Fig. 7E) has roundish cells with a large, basal nucleus and large, pale-rose secretory droplets. These cells occur together with small, ciliated cells that are broad at the base and taper towards the lumen. Their small nucleus stains deep-blue. The mucous gland appears blue to fuchsia-stained and reveals an epithelium with two cell types, namely columnar secretory cells with a granulated cytoplasm and a small nucleus in a basal position, and ciliated cells with a small nucleus (Fig. 7F).

Central nervous system and sensory organs

Anatomy. The CNS is situated near the intersection of the pharynx and the anterior part of the oesophagus, and forms a circumoesophageal ring (Fig. 8). A pair of very small buccal ganglia is located below the buccal mass. Close to the site of emergence of the rhinophores, there is a pair of small, pedunculate rhinophoral ganglia. The cerebral and pleural ganglia are fused to a pair of spheroid cerebropleural ganglia, which form the largest part of the CNS. They give rise to one main, paired nerve running towards the intestine (Fig. 8). The smaller paired pedal ganglia are located below the cerebropleural ganglia and the oesophagus. Three main nerves (Fig. 8) can be seen leaving these ganglia, all running into ventral direction.

The paired eyes are situated directly laterally from the cerebropleural ganglia, and are connected with the optic ganglia by the long optic nerves.

Histology. Ganglia are surrounded by a perineurium (Fig. 9A, B, E, F) that is formed by fibroblasts and contains some dispersed muscle cells. Neuronal perikarya are located in the ganglion peripheries, whereas axons run in the central neuropil. Many neuronal somata form axon bundles that leave the ganglia as commissures, connectives or nerves. The cerebropleural ganglia are interconnected by a thick cerebropleural commissure (Fig. 9A, B). In these ganglia we distinguish four categories of nerve cell bodies on the basis of their size: giant, large, medium-sized and small. The giant neurons were only observed in the cerebropleural ganglia. They measure ca. 70 μm in diameter, are spheroid, and have a large, central, blue nucleus (Fig. 9C). The large nerve cells (diameter ca. 40 μm), have a flattened apex that faces the perineurium (Fig. 9D). Like the medium-sized (diameter ca. 25 μm) and small neurons (diameter ca. 15 μm) they are present in all ganglia of the CNS. Many cells can be individually identified on the basis of their size, shape and their position in a ganglion, in each specimen studied.

The eyes are ovoid and have a largest diameter of ca. 60 μm. Their large, sea-blue-staining lens has a largest diameter of about 35 μm. The melanin at the inner face of the retina appears as small, deep-brown dots (Fig. 9E).

The paired statocysts are situated between the cerebropleural and pedal ganglia and reveal several otocoria in their lumen (Fig. 9F).
Egg mass

The light-cream coloured egg mass is a thin, undulating ribbon of about 5 mm length (Fig. 10A). It is filled with numerous tiny, oval, encapsulated eggs, each with a mean diameter of about 90 μm. The egg mass is deposited over hydroid and bryozoa colonies. After collecting an egg mass, eggs continued to develop, with one embryo per capsule (Fig. 10B).

Discussion

In this study we described in detail the anatomy, morphology and histology of the nudibranch, *D. uva* from the Chilean coast. The description clearly reveals all the diagnostic characters of the genus *Doto*, i.e., with respect to the body and foot shape, cerata, rhinophores, position of the anus and nephroproct, frontal veil, masticatory border, teeth, and penis (Thompson and Brown, 1984; for listing see Intro-
Each of these characters may not be unique for *Doto uva*, but it is the specific combination of these characters that separates *Doto uva* from other *Doto* species: a digestive system with bilobed salivary glands and a reproductive system characterized by (1) a voluminous ampulla, (2) a prostate gland consisting of two parts, (3) a short deferent duct, (4) an unarmed penis, (5) a seminal receptacle with internal lobes, (6) an oviduct with an internal valve, and (7) a vagina connected via a sphincter to the seminal receptacle.

Particular aspects of the characters typical for *D. uva* will be discussed below in a comparative context, as will new data on the animal’s morphology, anatomy and histology. Finally, we will consider if *D. uva* from the Chilean coast may be different from *D. uva* from the Brazilian coast as described by Marcus (1955, 1957).

### *Doto uva* from the Chilean coast

The peculiar globular shape of the voluminous salivary gland of *D. uva*, with large cells surrounding the gland, seems to be similar to that described for other *Doto* species, namely *D. coronata* (Gmelin, 1791) by Hecht (1896) and Brygider (1914), *D. pinnatifida* (Montagu, 1804) by Hecht (1896) and, more recently, *D. varians* MacFarland, 1966 by MacFarland (1966), which is synonymous with *D. kya* Marcus, 1961 (McDonald, 1983). The giant

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*Fig. 6. Histology of reproductive system. A: Ampulla (a). B: Prostate (pr) with globular cells (arrow) around the lumen (asterisk). C: Epithelium of the prostate with columnar cells (arrow) around the lumen (asterisk). D: Penial bulb (pb), unarmed penis (p) and vagina (asterisk). cg, capsule gland; d, vas deferens. Scale bars: 40 μm.*
cells in *D. uva* are similar in diameter to those of *D. varians* (diameter about 150 μm), but larger than those described for *D. coronata* and *D. pinnatifida*. The diversity of secretory cell types in salivary glands of molluscs is well-established (Boer et al., 1967; Moura et al., 2003) but the nature of the large...
surrounding cells in *D. uva* is obscure. In view of the granulated appearance of the cytoplasm, the cells may have a secretory function. Possibly, they possess a high degree of polyploidy, as is common for large-sized cells in molluscs in general (see e.g. Boer *et al.*, 1977).

Remarkably, Marcus (1957) reported that the *Doto* reproductive system lacks a seminal receptacle but has, instead, a voluminous bursa copulatrix (spermatheca) (Fig. 5B). Possibly, this description of a bursa is based on a misinterpretation, in view of the fact that also other Pacific and Atlantic *Doto* species lack a bursa copulatrix (O'Donoghue, 1921; Marcus, 1961), e.g. *D. columbiana* O'Donoghue, 1921, *D. amyra* Marcus, 1961 and *D. kya*. A similar arrangement of the reproductive system as found by us in *D. uva*, was reported for *D. columbiana* which also reveals a long vaginal duct. However, a difference is that in *D. columbiana* the sphincter is situated where the vagina opens into the female gland, and not near the connection site of the vagina with the seminal receptacle, as in our specimens.

Our study agrees with that of Marcus (1959) in that the reproductive system of *D. uva* from the Chilean coast contains one, very large and piriform seminal receptacle, but our specimens lack a bursa copulatrix whereas Marcus (1959) described a small one. However, as our serial section study strongly suggests, the structure Marcus described as bursa copulatrix most likely is the first part of the vaginal loop (Fig. 5A, C).

Taking the description of Marcus (1961) into account, Schrödl (2003) has suggested that *D. uva* and *D. kya* are closely related. Like *D. uva*, *D. kya* occurs at the Pacific coast, but should be restricted to North-America, living from Vancouver Island to California (Behrens, 1991). In the present study we observed several differences between the two species, e.g. with regard to body coloration pattern, the absence of dark rings on the cerata in *D. kya*, and the strongly ramifying pseudobranchs that hardly extend outside the cerata of *D. uva*. Therefore, we consider *D. uva* and *D. kya* as distinct species.

The other two species of *Doto* known from the Brazilian coast, *D. pita* Marcus, 1955 and *D. caramella* Marcus, 1957, are clearly different from *D. uva* in the pattern of coloration, shape and distribution of the cerata, tubercles, and shape of their teeth (Marcus, 1955, 1957, 1961).

This is the first detailed report on the anatomy and histology of the CNS of *D. uva*. We have distinguished four types of nerve cells on the basis of cell body size. These size differences may reflect different degrees of polyploidy that have been shown to be related to the amount of DNA content in the nuclei of nerve cells of pulmonate gastropods (Boer *et al.*, 1977). Obviously, this distinction is rather artificial as the functional diversity of neurons in the molluscan CNS is much higher than might be deduced from the presence of only four neuronal size classes. However, the distinction is functional in that it might form a basis to compare homologous neuron types between different nudibranchs and molluscan species in general. For instance, because of their position and flattened apical shape, the large nerve cells in the cerebropleural ganglia of *D. uva* may be homologous with the large neuromuscular growth hormone-producing cells in the cerebral ganglia of pulmonates (Geraerts, 1976; Roubos and van de Ven, 1987).

**Comparison of Doto uva from Chile with Doto uva from Brazil** (Table 1)

*D. uva* was described for the first time at the Brazil-
ian coast, by Marcus (1955, 1957). Later Marcus (1959) extended this description by studying *D. uva* at the coast of Chile, concluding that the specimens from both locations belong to the same species. The present data might challenge this conclusion, because our observations on the Chilean *D.*
uva and those of Marcus (1955, 1957) and more recently of ourselves on D. uva from the Brazilian coast, reveal large varieties in some characters among both specimens at both locations, raising the possibility two D. uva species occur mixed at both locations. In this respect, however, the following observations are relevant. First, irrespective of the location, it appears that the body coloration pattern of D. uva is white, yellow or pink, the spots dispersed over the body black or brown and the ramifications of the digestive gland pink, brown or greyish (Marcus, 1955, 1959; Schrödl, 1996; 2003; Schrödl et al., 2005; M.A. Fischer, unpublished results, present data; Table 1). Furthermore, cerata have either subapical dark-grey rings (Marcus, 1955, 1959), subapical black rings (present study) or basal black rings (Schrödl, 1996). Although these colour variations might be due to the existence of at least two species within a given population, it seems very likely that the differences are related to differences in individual developmental stage, feeding condition or to another physiological condition. A definite answer to this issue cannot be given as long as the biology of D. uva from the Pacific coast remains largely unknown.

Another source of variation within both the Chilean and Brazilian D. uva populations, is the number of radular teeth. We counted in two radula of D. uva from the Chilean coast, 61 and 63 teeth, which fits with Schrödl (2003), who counted 61-67 teeth in D. uva from Chile. Our specimens measured ca. 4.0 mm in preserved length, which is similar to the lengths of the specimens of D. uva investigated by Schrödl (2003) (4.5 and 5 mm). In contrast, Marcus (1955, 1957, 1959) described that the radula of D. uva from both Brazil and Chile has as many as 86 teeth. The holotype studied by Marcus (1955) measured 2 mm in preserved length, but unfortunately, the lengths of the other specimens studied by Marcus (1957, 1959) have not been reported. Consequently, the precise relationship between radular teeth number and body length remains to be established and, therefore, this number cannot be used as a reliable parameter to distinguish different species of D. uva.

A third character of interest when considering the issue of species distinction, concerns the rachidian tooth. In our specimens this tooth is horse-shoe-shaped, i.e., asymmetrical, with a central small cusp and 2-5 lateral denticles. However, Marcus (1955, 1959) described the rachidian tooth of D. uva from both Brazil and Chile to have only 2-3 lateral denticles whereas in his drawing of D. uva from Chile (Marcus, 1959) the tooth shows a central cusp with as many as 6 lateral denticles. Possibly, this variation in denticle number is related to the animal’s age, but it could, again, also point to different species existing within each of the two localities.

The scheme of the reproductive system made by Marcus (1957, 1959) shows some remarkable differences with our scheme, mainly in the shape of the vagina and position of the sphincter (Fig. 5A-C) that we attributed to misinterpretation of the sequence of the serial histological sections.

Lemche (1976) has proposed morphological and
biological criteria to distinguish between European *Doto* species, e.g. rhinophoral sheath shape, food preference and shape of the egg mass. The rhinophore seems to be frontally tapered and more protruded in specimens from Brazil, but comparing the preserved specimens from Marcus (1957) and Marcus (1959) there is not a big difference in the rhinophoral shape (Fig. 1D, F, G). As long as data on food and egg masses are not available for the Brazilian populations, these criteria cannot assist in solving the question as to the potential presence of two different species of *D. uva* from the South American coast.

Taken these arguments together, we propose to consider the Brazilian and Chilean populations of *D. uva* as belonging to the same species.

**Acknowledgements**

The authors wish to thank Mrs Elsa Gabriela Espinoza and Mr Ronaldo Rocco for assistance in collecting the material. We are also grateful to Mr Patricio Zabala (Pontificia Universidad Católica de Chile), Mr Anders Warén (Swedish Museum of Natural History) and Dr Luiz Ricardo Simone (Museu de Zoologia da Universidade de São Paulo) for making available specimens of *D. uva*. We also thank Dr Heike Wägele (University of Bonn, Germany) for providing laboratory facilities and advice, Dr Lucas Cervera Currado (University of Cádiz, Spain) for literature advise, Dr Elisabeth Pierson (Radboud University Nijmegen) for technical assistance, and Dr Victoriano Urgorri (University Santiago de Compostela) for comments on the manuscript.

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**Table 1. Comparison of characters and geographic distribution of *Doto uva* Marcus 1955 from the Chilean coast (present study) with other**

| References | Length (mm) | Coloration | Cerata/row |
|------------|-------------|------------|------------|
| Marcus, 1955 | 4.5 A; 2.0 P | White, transparent body with pale-orange internal organs; rose digestive gland; white cerata; black spots on tubercles, body dorsum and head | 5 |
| Marcus, 1957 | 8.0-9.0 A | One specimen with pale-rose body, another with pale-yellow body | 7-8 |
| Marcus, 1959 | 4.0-6.5 P | White body with dark spots on dorsum and one dark spot on the rhinophoral sheath; dark rings in tubercles | 3-8 |
| Schrödl, 1996 | until 8 A | White body with black spots; cerata with black rings on tubercle basis | 6 |
| Schrödl, 2003 | until 8 P | Whitish body with black spots | Up to 6 |
| Present study | 5-9 A 3-6 P | Yellow-hyaline body with brown spots on dorsum, head, lateral flanks, cerata, anal papilla and rhinophoral sheath; digestive gland appears as dorsal, brown or greyish band; tubercles with black ring; rhinophores with white small spots | 4-7 |
specimens of *Doto uva*. A, alive; P, preserved; -, not studied.

| Radula | Rachidean tooth | Vagina | Seminal receptacle | Geographic distribution |
|--------|-----------------|--------|---------------------|-------------------------|
| 86 × (0.1.0) | Central cup and 2-3 lateral denticles | - | - | Sao Sebastiao Island, Sao Paulo, Brazil |
| 86 × (0.1.0) | - | Short, wide and muscular | Large and piriform, described aspermatotheca | Ihiabela and Sao Paulo, Brazil |
| 86 × (0.1.0) | Central cup and 2-3 lateral denticles | Short; bursa type | Large and piriform | Golf Ancud, Chile (41°46’30″S; 73°06’45″W) |
| - | - | - | - | Columo Bay, Chile (36°32’S; 72°57’W) |
| 61-71 × (0.1.0) | Central cup and 2-5 lateral denticles | - | Stalked allosperm receptacle | Columo Bay, Chile |
| 61-63 × (0.1.0) | Central cup and 2-5 lateral denticles | Long; loop at the upper part of penial bulb. Bulbous close to seminal receptacle | Large and piriform with internal lobes | Punta Blanca, Tocopilla, Chile (22°04’S; 70°12’W); La Herradura, Chile (29°58’S; 71°22’W); Tongoy, (30°15’S; 71°30’W) |

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Received: 3 February 2006
Accepted: 28 June 2006
