

## The earliest fossil record of the poorly known family Condylocardiidae from Argentina

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**ABSTRACT.** The scarcely known family Condylocardiidae (Bivalvia: Archiheterodonta) is poorly represented in the fossil record and their living representatives are also poorly known. This work presents a new representative of the family from the early Pliocene of marine terrace of Cerro Laciár (Santa Cruz Province). *Carditella pitufina* sp. nov. is described and characterized by a shell large for the genus, 15 radial ribs as wide as interspaces, high hinge plate and broad and large hinge teeth. The poor record of the family may be due to an identification bias, probably because the tiny size of specimens and its similarity with carditids. This new species resembles to the living *Carditella tegulata* from southern Argentina and Chile and they could be closely related. *Carditella pitufina* sp. nov. represents the most ancient record of *Carditella* in South America and the most ancient record of Condylocardiidae in Argentina.

*Keywords:* Archiheterodonta, Bivalvia, Carditella, Condylocardiidae, Patagonia, Pliocene.

**RESUMEN.** El registro fósil más antiguo de la poco conocida familia Condylocardiidae en Argentina. La muy poco conocida familia Condylocardiidae (Bivalvia: Archiheterodonta) está pobremente representada en el registro fósil y las especies vivientes del grupo han sido poco estudiadas. Este trabajo presenta un nuevo representante de la familia, procedente de la terraza marina de cerro Laciár (Provincia de Santa Cruz), de edad Plioceno temprano. *Carditella pitufina* sp. nov. es descrita y caracterizada por presentar valvas grandes para el género, 15 costillas radiales tan anchas como sus interespacios, charnela alta y ancha y dientes charnelares grandes. El pobre registro de la familia es referido a un sesgo de identificación, debido probablemente al pequeño tamaño de los especímenes y a su similitud con los cardítidos. Esta nueva especie se asemeja a la especie viviente *Carditella tegulata* del extremo sur de Argentina y Chile y podrían estar estrechamente emparentadas. *Carditella pitufina* sp. nov. representa el registro más antiguo del género *Carditella* en América del Sur y el más antiguo de un Condylocardiidae para Argentina.

*Palabras clave:* Archiheterodonta, Bivalvia, Carditella, Condylocardiidae, Patagonia, Plioceno.

## 1. Introduction

The poorly known family Condylacardiidae Bernard, 1896 is characterized by its very small size and narrow hinge with anterior and posterior lateral teeth in each valve (according to Middelfart, 2002a). Major aspects of the biology of condylacardiids are unknown (Middelfart, 2002a). Their tiny size and relative low abundance may be the reason of the scarce knowledge and recognition of living specimens. The family was related to the Carditidae and placed together in the superfamily Carditoidea (Dall, 1903a; Thiele, 1935) and recently was considered a member of the clade Archiheterodonta (Giribet, 2008). Posterior analyses of molecular sequences placed Condylacardiidae as a clade nested within Carditidae (González and Giribet, 2014). This placement agrees with the proposal of Bernard (1897) who considered the condylacardiids derived by neoteny from carditids, and Coan (2003) who suggested a polyphyletic origin for the group.

The Condylacardiidae are frequently distinguished from other archiheterodonts by their tiny size, lateral teeth in both valves, presence of internal ligament, and the presence of large prodissoconchs (Middelfart, 2002b). According other authors (Huber, 2010; Güller and Zelaya, 2013), some genera of condylacardiids present both, internal and external ligament (or exclusively external) and small prodissoconchs. These discrepancies lead to discussions about the taxonomic composition of the family (Middelfart, 2002b; Coan, 2003; Huber, 2010; Coan and Valentich-Scott, 2012).

Living Condylacardiidae were reported from Australia (Middelfart, 2002a, b), tropical eastern Pacific (Coan, 2003), eastern Atlantic (Salas and Rolán, 1990) and southern Atlantic and Pacific (Güller and Zelaya, 2013). Fossil Condylacardiidae were reported from the Paleocene of Nigeria (Adegoke, 1977); Eocene of France (Sanders *et al.*, 2015), United States (Palmer and Brann, 1965), and New Zealand (Maxwell, 1992); Oligocene of United States (Dockery, 1982); Miocene of Australia (Tate, 1886; Ludbrook, 1953), New Zealand (Gage, 1957), Belgium (Louwey *et al.*, 2010), Germany (Moths *et al.*, 2010), and Indonesia (Beets, 1983); Pliocene of New Zealand (Laws, 1940), Belgium (Marquet, 2002), Trinidad and Tobago (Jung, 1969), Indonesia (Oostingh, 1935) and United States (Ward and Blackwelder, 1987); and Quaternary of South Africa (Kensley and Pether, 1986) and Costa Rica (Olsson, 1942). A possible previous record comes from the

Cretaceous of North America (Stephenson, 1941) represented by the dubious species *Uddenia texana* Stephenson, 1941.

There are only three fossil records of the Condylacardiidae in South America. The first is founded in the late Miocene of Uruguay (Camacho Formation) and refers to *Warrana antiqua* (Figueiras, 1990) (formerly described as *Americuna antiqua*). The second and third correspond to two mentions listed but not illustrated by Gordillo *et al.* (2005) from middle Holocene of Tierra del Fuego (Argentina) -*Carditella naviformis* (Reeve, 1843) and *Carditopsis flabellum* (Reeve, 1843)-. Nowadays four species of living condylacardiids are mentioned in the region, from southern Peru, Chile and Tierra del Fuego (Güller and Zelaya, 2013). The aim of this contribution is to report and to describe the most ancient fossil record of Condylacardiidae of Argentina, represented by a new species from early Pliocene marine terraces of Santa Cruz (Argentina).

## 2. Geological setting

Fossil locality of Terrace of Cerro Laciari (Fig. 1), studied by Feruglio (1933, 1950) and assigned to his System Terrace I, are located 50 km west to Puerto Deseado, Santa Cruz Province. These exposures are placed at 170-185 meters above sea level, overlaying the marine Monte León Formation (early Miocene) and consist of few meters of loose gravel with calcareous sandy matrix (Fig. 1). Condylacardiids are founded within the bivalves *Scalariocardita laciarina* (Feruglio, 1954), *Kolmeris tehuelchana* (Ihering, 1907), *Ostrea ferrarisi* d'Orbigny, 1842, "*Chlamys*" *actinodes* (Sowerby, 1846), and the gastropods "*Trophon*" *varians* (d'Orbigny, 1842) and *Pachycymbiola feruglioi* (Doello Jurado, 1931). The age of these marine fossiliferous beds has been discussed, and assigned to Miocene (Ameghino, 1906), Pliocene (Feruglio, 1950, 1954) or Pleistocene (Feruglio, 1933). Recently, Del Río *et al.* (2013) indicated for these deposits an early Pliocene age ( $5.10 \pm 0.21$  Ma, Zanclean) based on  $^{87}\text{Sr}/^{86}\text{Sr}$  analysis from valves of the pectinid "*Chlamys*" *actinodes*.

## 3. Material and methods

The fossil and recent specimens used for comparisons are housed at Museo Argentino de Ciencias Naturales "Bernardino Rivadavia". Shells

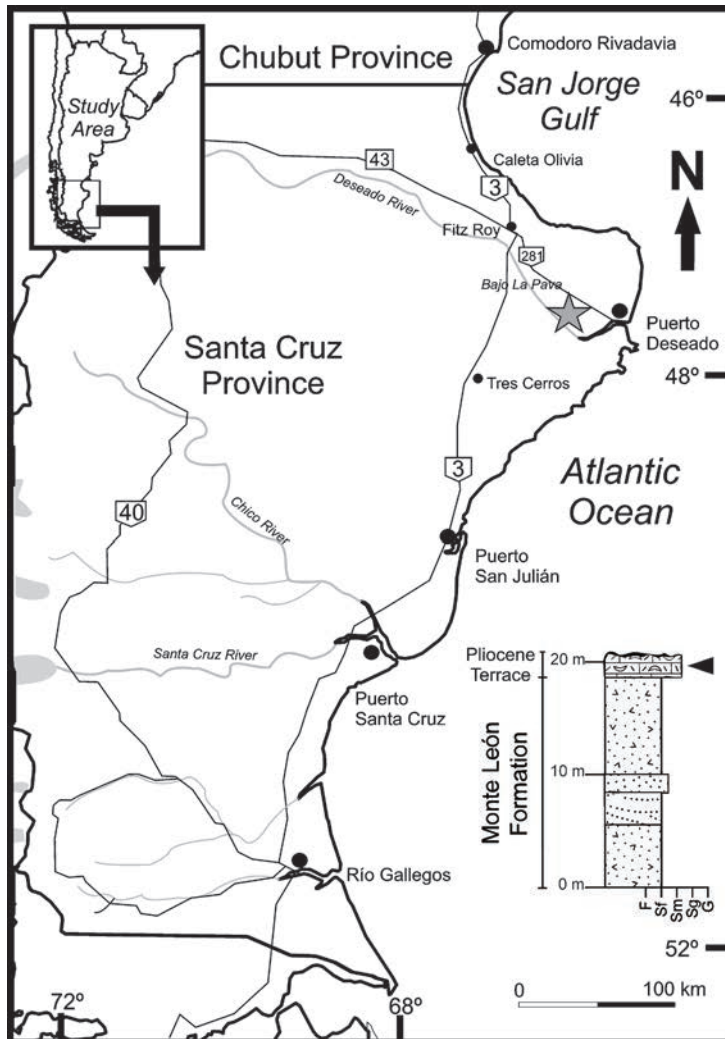


FIG. 1. Geographical provenance (indicated by a grey star) of studied fossils and lithological section at Terrace of Cerro Laciar.

were viewed with a Leika binocular magnifying glass. The preservation of fossil shells is moderate, they shows some degree of surface and edge alteration, and bioerosion product of the action of boring sponges *-Entobia-* (Gordillo *et al.*, 2014). Besides, this attributes did not difficult the identification of morphological features. Hinge terminology follows Lamy (1917), other morphological terms (*e.g.*, “ligamental pit”) follows Middelfart (2002a, b).

**Institutional abbreviations.** MACN-In (Invertebrates Collection of the Museo Argentino de Ciencias Naturales “Bernardino Rivadavia”, Buenos Aires, Argentina); MACN-Pi (Paleoinvertebrates Collection of the Museo Argentino de Ciencias

Naturales “Bernardino Rivadavia”, Buenos Aires, Argentina).

#### 4. Systematic paleontology

**Order Archiheterodonta**  
**Family Condyllocardiidae Bernard, 1896**  
**Subfamily Carditellinae**  
**Kuroda, Habe and Obaya, 1971**

**Remarks.** Chavan (1969) and Middelfart (2002a, b) restricted the family for species with an external ligament only and divided the Condyllocardiidae in two subfamilies: Condyllocardiinae Bernard, 1896 and

Cuninae Chavan, 1969. They placed small species with external and internal (*Carditella* Smith, 1881) or only external ligament (*Carditellona* Iredale, 1936 and *Carditellopsis* Iredale, 1936) within the family Carditidae, despite other characters. This approach was not followed by Coan (2003) who considered *Carditella* as a condylocardiid. Kuroda *et al.* (1971) suggested a third subfamily, Carditellinae, for these closely related species, following the original concept of Iredale and McMichael (1962) when they grouped the genera *Carditellona* and *Carditellopsis*. This approach with three subfamilies instead two was accepted by Huber (2010) and is followed here. Carditellinae taxa share with the Cuninae the presence of a small and poorly defined cup-shaped prodissoconch and triangular-shaped right middle and left posterior teeth.

### Genus *Carditella* Smith, 1881

**Type species.** *Carditella pallida* Smith, 1881 by subsequent designation (Dall, 1903b) (Recent, southern Chile) (junior synonym of *Carditella tegulata* according to Güller and Zelaya, 2013).

**Emended diagnosis.** Shell small and flabellate. Lunule elongated. External sculpture of 11-15 radial ribs. Hinge composed by two left cardinal and three right cardinal teeth. Each valve also has two lateral teeth, one nearly marginal on the one side, the other on the opposite side being well within the outer edge, with a groove between it and the anterior margin. External and internal ligament small, internal placed beneath the apex of the valves in a small resilifer (Modified from Smith, 1881).

**Remarks.** In its original description, Smith (1881) recognized that external sculpture resembles carditids. The genus *Carditella* is characterized by their flabellate shell-outline with wide radial ribs covered by subrectangular nodes, hinge with two left and three right cardinal teeth and anterior and posterior lateral teeth in both valves. Also, *Carditella* has external and internal ligament, the latter marked by a ligamental pit beneath the apex in left valve. The presence of strongly developed lateral teeth and a small ligamental pit allow to place the new species in the condylocardiid genus *Carditella*. Hinge configuration and external sculpture do not differ from other species of *Carditella*.

Two fossil species of *Carditella* were recorded: *C. baloguni* Adegoke, 1977 (late Paleocene, Nigeria)

and *C. aldrichi* (Casey, 1903) (middle Eocene-early Oligocene, United States of America). *Carditella calipsamma* Carrington and Kensley, 1969 (late Pliocene-early Pleistocene, South Africa) seems to be a representative of the genus *Carditellona* Iredale, 1936 instead a southern African *Carditella* species. Nowadays, the genus *Carditella* is represented by *C. naviformis* (Reeve, 1843) (southern South America), *C. tegulata* (Reeve, 1843) (southern Peru to Tierra del Fuego), *C. semen* (Reeve, 1843) (southern Peru to Chile), *C. exulata* Smith, 1885 (Tristan da Cunha), *C. galapagana* Coan, 2003 (western coast of Colombia and Galapagos Islands), *C. marieta* Coan, 2003 (western coast of Mexico), *C. capensis* Smith, 1885 (Recent, Angola and South Africa), *C. delli* Crozier, 1966 (northern New Zealand), *C. iejimensis* Hayami and Kase, 1993 (southern Japan), *C. laticosta* Smith, 1904 (South Africa), *C. mawsoni* Dell, 1972 (Weddel Sea), and *C. tridacnula* Oliver and Chesney, 1997 (Oman).

### *Carditella pitufina* sp. nov.

Fig. 2.1-13, 3.1

**Etymology.** The specific epithet refers to the Belgian comic characters created by Peyo, “Les Schtroumpfs” (called “The Smurfs” in English and “Los Pitufos” in Spanish speaker countries). These creatures are characterized by their tiny sizes as the condylocardiids bivalves. This epithet is expressed in Spanish diminutive form to emphasize the reference.

**Type material.** Holotype MACN-Pi 6344 (a left valve). Paratype MACN-Pi 6344 (a left and three right valves).

**Type locality and horizon.** Terrace of Cerro Laciár, Santa Cruz Province (lower Pliocene marine terraces).

**Diagnosis.** Shell large for the genus (length range 6.27-7.12 mm), external sculpture of 15 radial ribs as wide as interspaces, hinge plate high and hinge teeth broad and large.

**Description.** Shell large-sized for the genus (length range 6.27-7.12 mm) (Table 1), mainly subtriangular; convex anterior and posterior margins, rounded ventral margin, dorsal margin sharply sloping and very curved, angle between dorsal and posterior margins smooth, angle between dorsal and anterior margins weak marked. Umbo small, pointed or slightly rounded, anteriorly directed, placed at centre to anterior third of the valve length. Lunule large,

elongated and narrow, slightly concave, without sculpture and smoothly separated from remaining surface of shell. Escutcheon large and elongated, smooth, larger than lunule. Prodissoconch small, eroded in all specimens studied. Hinge plate high, three cardinal and two lateral teeth in right valve and two cardinal and two lateral teeth in left valve. Right valve hinge with rounded ventral edge; anterior cardinal tooth thin and straight, anteriorly inclined; middle cardinal tooth subtriangular, slightly inclined posteriorly, with broad base; posterior cardinal tooth thin, long and straight; anterior lateral tooth elongated, wide, separated from lunular margin; posterior lateral tooth elongated, high, not differentiated from anterior

margin. Left valve hinge with a straight ventral edge; anterior cardinal tooth curved, anteriorly inclined, with broad base; posterior cardinal tooth long and curved, inclined posteriorly, with broad base and acute apex; anterior lateral tooth thin and high, close to anterior margin; posterior lateral tooth long and wider than anterior lateral. Small ligamental pit placed below beak in left hinge. External sculpture of 15 radial ribs as wide as interspaces, with subrectangular transverse section, covered by rectangular nodes; interspaces with subrectangular transverse section. Pallial line at a quarter of total valve height. Inner ventral margin crenulated; crenulations subrectangular and truncated, covering the entire margin.

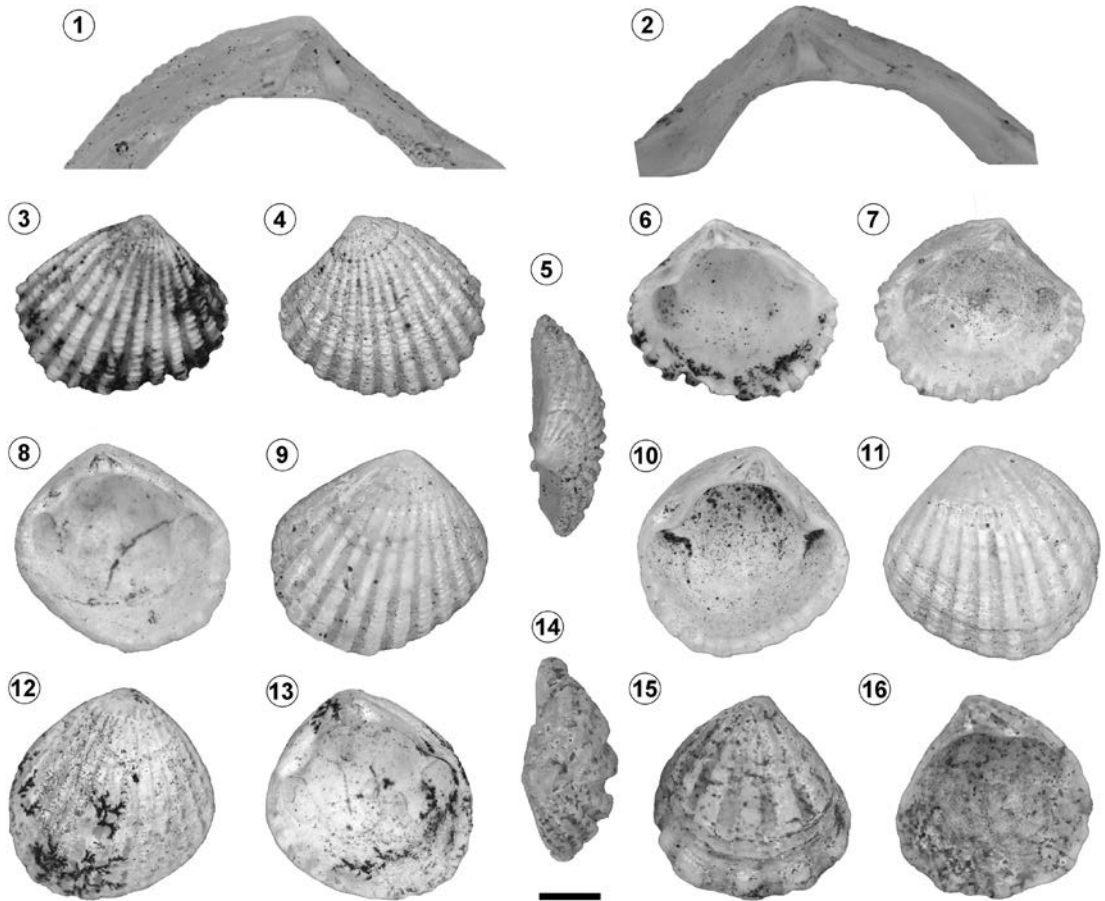


FIG. 2. 1-13. *Carditella pitufina* n. sp., MACN-Pi 6344; 1. Left hinge (Holotype); 2. Right hinge (Paratype); 3. Right valve, lateral view (Paratype); 4. Left valve, lateral view (Holotype); 5. Left valve, dorsal view (Holotype); 6. Right valve, internal view (Paratype); 7. Left valve, internal view (Holotype); 8. Right valve, internal view (Paratype); 9. Right valve, lateral view (Paratype); 10. Left valve, internal view (Paratype); 11. Left valve, lateral view (Paratype); 12. Right valve, lateral view (Paratype); 13. Right valve, internal view (Paratype); 14-16. *Kolmeris tehuelchana* (Ihering, 1907), MACN-Pi 2214; 14. Right valve, dorsal view; 15. Right valve, lateral view; 16. Right valve, internal view. Scale bar represents 5 mm for 1-2, and 2 mm for 3-16.

TABLE 1. MEASUREMENTS OF STUDIED SPECIMENS OF *CARDITELLA PITUFINA* SP. NOV.

Specimen	Length (mm)	Height (mm)
MACN-Pi 6344-1 (Paratype)	7.04	6.8
MACN-Pi 6344-2 (Paratype)	6.27	6.03
MACN-Pi 6344-3 (Paratype)	6.91	6.79
MACN-Pi 6344-4 (Paratype)	6.63	6.44
MACN-Pi 6344-5 (Paratype)	6.8	7.01
MACN-Pi 6344-6 (Holotype)	6.85	6.06
MACN-Pi 6344-7 (Paratype)	7.12	6.17

**Remarks.** Shell of the new species shows some variability from more to less triangular outlines. Placement of umbos is more or less centrally. Development of shell margins and radial ribs varying in studied specimens but they are due to taphonomic processes (e.g., surface alteration).

This new species strongly resembles to *Carditella tegulata* (Reeve, 1843) (Recent, Valparaíso [33° S] to Beagle Channel [55° S]) (Güller and Zelaya, 2013: Fig. 5) but is distinguished by its quite larger shell with narrower radial ribs, higher hinge plates, larger and broader cardinal teeth, and right hinge with less curved middle tooth. Also, the dorsal margin in the fossil species is more curved and the prodissoconch is smaller than those of *C. tegulata*. On the other hand, the new species can be distinguished from *C. baloguni* Adegoke, 1977 (Ewekoro Formation, late Paleocene, Nigeria) (Adegoke, 1977: pl. 50, Figs. 1-7) by its less prominent umbos, broader hinge and less numerous radial ribs without comarginal nodes.

*Carditella pitufina* sp. nov. resembles *C. aldrichi* (Casey, 1903) (Gospport Sand, Mint Spring and Byram formations, middle Eocene-early Oligocene, United States of America) (Dockery, 1982; pl. 24, figs. 12-13, pl. 25, figs. 1-5, 7) but the Argentinean species has a less elongated shell, less rounded posterior margin, and broader hinge with less divergent and more robust teeth.

*Carditella pitufina* sp. nov. can be separated from *C. naviformis* (Reeve, 1843) (Recent, Valparaíso [33°S] to Malvinas Islands [51° S]) (Güller and Zelaya, 2013, fig. 6) and *C. semen* (Reeve, 1843) (Recent, Isla Lobos de Afuera, Peru [7° S] to Isla Blanca, Chile [27° S]) (Güller and Zelaya, 2013:

Fig. 7) because the expanded outlines in the living species, with the presence of a more prominent prodissoconch separated from teleoconch by a bulging rim, fine comarginal lines between radial ribs, and with smaller and narrower hinge teeth.

## 5. Discussion

There are very few previous records of the family Condyllocardiidae in South America (Figueiras, 1990; Gordillo *et al.*, 2005), which may be due to an identification bias. Shells of this group are very small sized (see Table 1 for *Carditella pitufina* sp. nov. sizes), and hinge features are visible only by magnification. Small bivalves are difficult to assign taxonomically (Valentine *et al.*, 2006). Externally, condyllocardiids resemble to small specimens of Carditidae taxa, and Smith (1881) already mentioned the external similarity between *Carditella* and carditids. At the marine terraces of Cerro Laciár, two small carditid species are recorded: *Kolmeris tehuelchana* (length range of 6.3-15.9 mm) (Fig. 2.14-16, Fig. 3.2, 3.4) and *Scalaricardita laciarina* (length range of 8.4-26 mm) (Fig. 3.3), but differences on adult size among them are noticeable (Fig. 3). Carditid species share external sculpture of radial ribs with *Carditella pitufina* sp. nov., but they present 25-27 (*S. laciarina*) or 13-15 ribs stacked at sides (*K. tehuelchana*) (Pérez and Del Río, 2017), while the condyllocardiid has 15 not stacked radial ribs. In *S. laciarina*, radial ribs are covered by subrectangular nodes as in *C. pitufina* sp. nov., but in *K. tehuelchana*, radial ribs are smooth (Pérez and Del Río, 2017). Also, carditids have not lateral teeth or ligamental pit. These features distinguish the new species from the known carditids of the Cerro Laciár marine terraces. Some juvenile specimens of *K. tehuelchana* with similar size of adult *C. pitufina* sp. nov. present these strong differences in hinge teeth and external sculpture (Fig. 2.15). Specimens of *C. pitufina* sp. nov. here described were founded among very small Carditidae specimens originally collected by E. Feruglio.

*Pleuromeris* Conrad, 1867 is another small carditid genus recorded in the late Oligocene-middle Miocene of Argentina (San Julián, Monte León and Carmen Silva formations). This genus is represented by two species, *P. sulcolumularis* (Ihering, 1907) and *P. fueguina* (Steinmann and Wilckens, 1908). *Pleuromeris* is characterized by small (less than

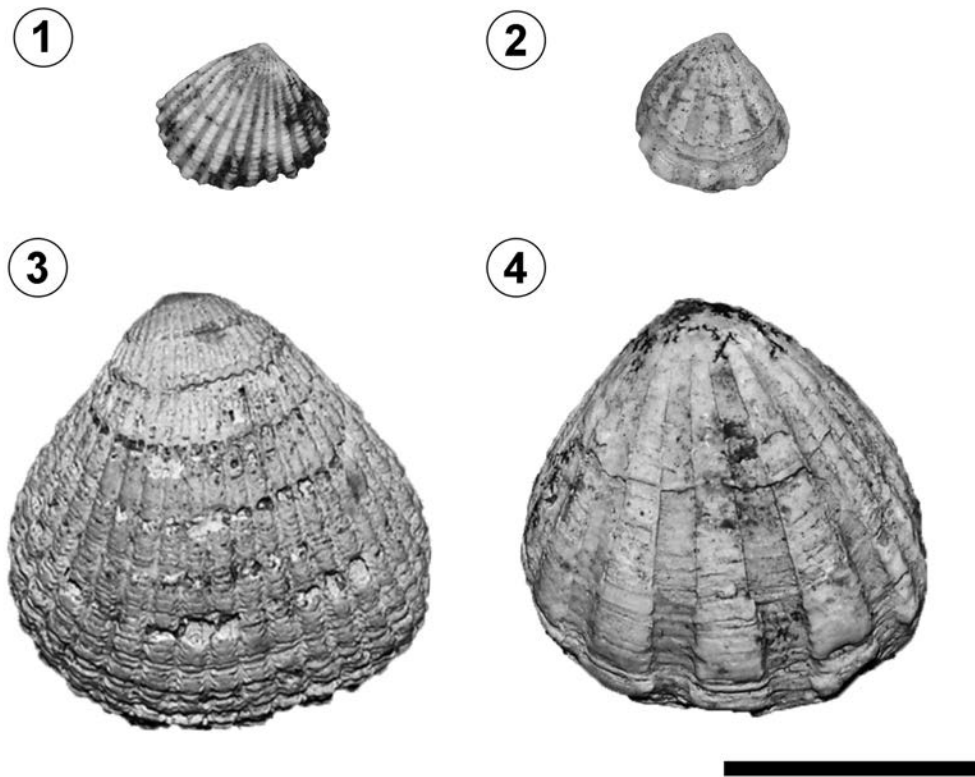


FIG. 3. Comparison between *Carditella pitufina* n. sp. and carditids of Terrace of Cerro Laciár. 1. *Carditella pitufina* n. sp., MACN-Pi 6344, right valve, lateral view (Paratype); 2. Juvenile *Kolmeris tehuelchana* (Ihering, 1907), MACN-Pi 2214, right valve, lateral view; 3. Adult *Scalaricardita laciarina* (Feruglio, 1954), MACN-Pi 2213, left valve, lateral view; 4. Adult *Kolmeris tehuelchana* (Ihering, 1907), MACN-Pi 361, left valve, lateral view. Scale bar represents 10 mm.

30 mm in length) and subtriangular outlined shells. This taxon has a broad and triangular right middle tooth and a reduced left anterior tooth that are not present in *Carditella*. Also, the condylocardiid genus has lateral teeth in both hinges. Some small species of *Pleuromeris* from New Zealand (e.g., *P. benthicola*, *P. ultima*) externally resemble to *Carditella* but they have wider radial ribs and a broader right middle tooth. Argentinian species of *Pleuromeris* have larger shells (near to 20-25 mm), broad right middle tooth and a higher number of radial ribs.

*Carditella* was present in the Paleocene of Africa and in the Oligocene-Eocene of North America, but this is its first Pliocene record. The present distribution of *Carditella* ranges from southern South America towards Pacific coast to Colombia and Mexico, and by Atlantic coast towards southern Africa. The genus is also present in New Zealand, Japan and Oman.

The species of Condylocardiidae from the southern South America are distributed from Tierra del Fuego towards Chile and Peru, and they are represented by four species: *Carditella tegulata* (Reeve, 1843), *C. naviformis* (Reeve, 1843), *C. semen* (Reeve, 1843) and *Carditopsis flabellum* (Reeve, 1843) (Güller and Zelaya, 2013). The new record of *C. pitufina* sp. nov. indicates a more northern distribution of the group in western Atlantic for Pliocene times, and the most ancient record of the genus in South America. *Carditella pitufina* sp. nov. is very similar to the Recent *C. tegulata*, which is recorded from middle Holocene (Gordillo *et al.*, 2005), and could be a Pliocene relative for this species.

Pliocene marine terraces of Argentina were recently recognized because their debated age (Del Río *et al.*, 2013), the faunal composition of Terrace of Cerro Laciár was not revised after Feruglio (1933,

1950, 1954), notwithstanding some mentions (Del Río and Martínez, 2006; Aguirre *et al.*, 2008; Pérez and Del Río, 2017; Trovant *et al.*, 2017). In contrast of Miocene molluscs of Patagonia (Martínez and Del Río, 2002; Del Río *et al.*, 2010), Pliocene molluscs seems to be more related to actual malacological faunal composition. According to the faunal list provided by Feruglio (1950) (and the previously mentioned posterior revisions), Terrace of Cerro Laciár includes a certain amount of extant taxa [*e.g.*, *Maetra isabelleana* d'Orbigny, 1846, *Mytilus edulis* Linnaeus, 1758, *Perumytilus purpuratus* (Lamarck, 1819), *Tegula patagonica* (d'Orbigny, 1835), *Trochita pileolus* (d'Orbigny, 1842), *Falsilunatia patagonica* (Philippi, 1845), *Trophon geversianus* (Pallas, 1774), “*Trophon*” *varians* (d'Orbigny, 1842), *Boreoscala magellanica* (Philippi, 1845)] and other taxa closely related to extant species [*e.g.*, *Amiantis laziarina* (Ihering, 1903), “*Chione*” *laciarina* Feruglio, 1954, *Urosalpinx* sp., *Pachycymbiola feruglioi* (Doello Jurado, 1931)]. The new record of *Carditella* are placed in the second group, and enriches the connections between Pliocene and modern faunas. New revisions of this mollusc fauna are needed for enlighten the origins of modern faunal composition of southern South America.

## 6. Conclusions

A new species of the genus *Carditella* Smith, 1881, *C. pitufina* sp. nov., is described from the early Pliocene marine terraces of Cerro Laciár, Santa Cruz Province, Argentina. This finding represents the most ancient fossil record of Condylacardiidae of Argentina and the most ancient record of the genus *Carditella* for the south end of South America.

## Acknowledgements

Thanks to the curators M. Longobucco (MACN-Pi), C. Del Río (MACN-Pi) and A. Tablado (MACN-In). Thanks to M. Belén Santelli (MACN-Pi) who suggested the idea of the specific name for the new taxa and M. Álvarez (MACN-Pi) for his suggestions. I am very grateful to P. Hanisch (MACN) for her help with the photography of very small shells. Also thank to M. Güller (MACN) for helping in the very preliminary stage of this research and the workers of the Library of MACN for their valuable assistance.

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