



This work is licensed under a Creative Commons Attribution License (CC BY 4.0).

Research article

[urn:lsid:zoobank.org:pub:2E0FAA1D-DA9A-4486-805F-9DA3DF928539](https://zoobank.org/pub:2E0FAA1D-DA9A-4486-805F-9DA3DF928539)

Taxonomy and distribution of Pectinariidae (Annelida) from Iceland with a comparative analysis of uncinal morphology

Julio PARAPAR^{1,*}, Verónica PALOMANES²,
Gudmundur V. HELGASON³ & Juan MOREIRA⁴

^{1,2}Departamento de Biología, Universidade da Coruña, 15008 A Coruña, Spain.

³Deceased 9 May 2020. Former address: RORUM ehf., Brynjólfsgötu 5, 107 Reykjavík, Iceland.

⁴Departamento de Biología (Zoología), Universidad Autónoma de Madrid, Cantoblanco, 28049 Madrid, Spain.

⁴Centro de Investigación en Biodiversidad y Cambio Global (CIBC-UAM), Universidad Autónoma de Madrid, 28049 Madrid, Spain.

*Corresponding author: julio.parapar@udc.es

²Email: veronica.palomanes@udc.es

⁴Email: juan.moreira@uam.es

¹[urn:lsid:zoobank.org:author:CE188F30-C9B0-44B1-8098-402D2A2F9BA5](https://zoobank.org/author:CE188F30-C9B0-44B1-8098-402D2A2F9BA5)

²[urn:lsid:zoobank.org:author:6C644341-D35B-42B6-9857-5F119457A424](https://zoobank.org/author:6C644341-D35B-42B6-9857-5F119457A424)

³[urn:lsid:zoobank.org:author:32B3520E-1D49-4B77-BF81-2AAE3FE76363](https://zoobank.org/author:32B3520E-1D49-4B77-BF81-2AAE3FE76363)

⁴[urn:lsid:zoobank.org:author:B1E38B9B-7751-46E0-BEFD-7C77F7BBBEF0](https://zoobank.org/author:B1E38B9B-7751-46E0-BEFD-7C77F7BBBEF0)

This paper is dedicated to Guðmundur Vidir Helgason who passed away on 9 May 2020, just before publication of this paper. Project Manager at RORUM, an environmental research and consulting company, he was previously a Project Coordinator for the BIOICE program (Benthic Invertebrates of Icelandic Waters) and Director of the Sandgerði Marine Centre from 1992 to 2013, being one of the organizers of the 7th International Polychaete Conference (Reykjavík, July 2001).

Abstract. Based on samples collected during the BIOICE project off Iceland, four species of marine annelids belonging to the family Pectinariidae were identified: *Amphictene auricoma* (O.F. Müller, 1776), *Cistenides granulata* (Linnaeus, 1767), *Cistenides hyperborea* Malmgren, 1865 and *Lagis koreni* Malmgren, 1866. Taxonomic remarks and data on geographical and bathymetric distribution are presented. The distribution of each species off Iceland was evaluated and two patterns were defined: *C. granulata* and *C. hyperborea* were mainly found in waters off the northeast coast, while *A. auricoma* and *L. koreni* were found on the southern coast. Several body characters with taxonomic value in this family were reviewed under the stereo microscope and scanning electron microscope, with special emphasis on the neuropodial uncini. Remarks on these special chaetae are included in the diagnoses.

Keywords. SEM, Polychaeta, distribution, species diversity.

Parapar J., Palomanes V., Helgason G.V. & Moreira J. 2020. Taxonomy and distribution of Pectinariidae (Annelida) from Iceland with a comparative analysis of uncinial morphology. *European Journal of Taxonomy* 666: 1–32. <https://doi.org/10.5852/ejt.2020.666>

Introduction

The BIOICE (Benthic Invertebrates of Icelandic Waters) expeditions started in 1992 aiming at characterizing the marine benthic fauna living in the 200-mile economic zone of Iceland in a depth range of 20 to 3500 m. Sampling was carried out at both sides of the Greenland–Iceland–Faeroe Ridge (GIF Ridge) that separates water masses with different oceanographic features (Hansen & Østerhus 2000). In this sense, several water masses associated with submarine ridges are present around Iceland, influencing the flow of several cold and warm water currents (Silva *et al.* 2017), dividing its coast into three distinct regions (Fig. 1A), and therefore limiting the distribution of the marine species (Brix & Svavarsson 2010; Meißner *et al.* 2014; Silva *et al.* 2017).

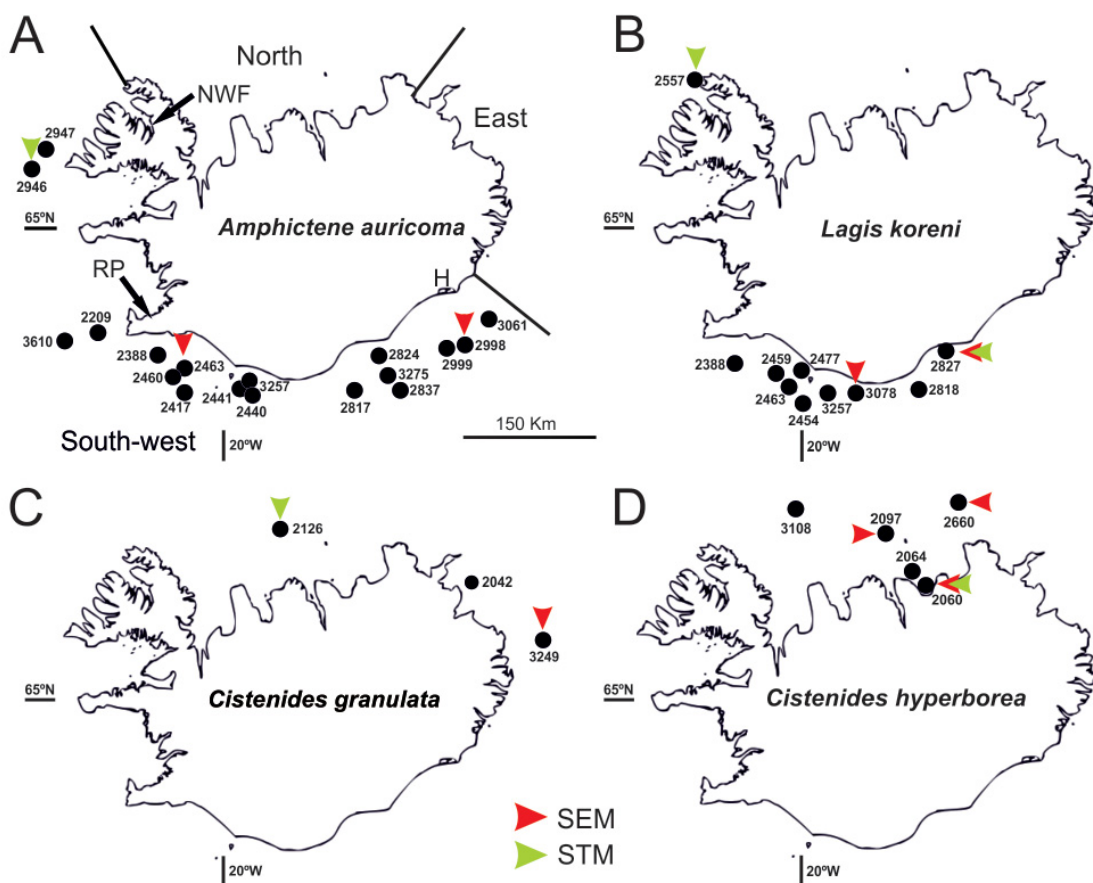


Fig. 1. Maps of Iceland showing distribution of the species of Pectinariidae found in BIOICE samples. **A.** *Amphictene auricoma* (O.F. Müller, 1776). **B.** *Lagis koreni* Malmgren, 1866. **C.** *Cistenides granulata* (Linnaeus, 1767). **D.** *Cistenides hyperborea* Malmgren, 1866. The three main coastal sectors around Iceland are shown in A. The specimens photographed and studied under SEM are indicated with green and red arrowheads, respectively. Abbreviations: H = Höfn; NWF = North-western fjords; RP = Reykjanes Peninsula.

The polychaete fauna (Annelida Lamarck, 1809) yielded by the BIOICE expeditions has been extensively studied in the last two decades, for instance by Petersen (2000: Fauveliopsidae Hartman, 1971), Kirkegaard (2001: Glyceridae Grube, 1850 and Goniadidae Kinberg, 1866), Sanfilippo (2001: Serpulidae Rafinesque, 1815), Sigvaldadóttir (2002: Spionidae Grube, 1850), Chambers & Woodham (2003: Cirratulidae Ryckholt, 1851), Parapar (2003, 2006: Oweniidae Rioja, 1917), Moreira & Parapar (2012, 2015: Sphaerodoridae Malmgren, 1867), Parapar *et al.* (2011a, 2011b, 2011c, 2013a, 2013b, 2014a, 2014b: Trichobranchidae Malmgren, 1866, Opheliidae Malmgren, 1867, Ampharetidae Malmgren, 1866 and Longosomatidae Hartman, 1944) and Zamora *et al.* (2020: Fauveliopsidae). However, other families well represented in such samples have not been studied yet such as the Pectinariidae Quatrefages, 1866.

The family Pectinariidae was traditionally composed of five genera (Malmgren 1866): *Pectinaria* Savigny in Lamarck, 1818, *Amphictene* Lamarck, 1818, *Cistenides* Malmgren, 1866, *Lagis* Malmgren, 1866 and *Petta* Malmgren, 1866; this classification was followed, for instance, by Long (1973), Fauchald (1977) and Wolf (1984). However, other authors only recognized *Pectinaria* and *Petta*, considering the other three as subgenera of *Pectinaria* (e.g., Hessle 1917; Fauvel 1927; Hartman 1941; Day 1967; Holthe 1986). We follow Hutchings & Peart (2002), who recognized the five aforementioned genera and provided a diagnosis for each genus, together with a table listing the major diagnostic characters; this systematic arrangement was also followed by recent workers such as Garraffoni & Camargo (2006), Nogueira *et al.* (2019) and Read & Fauchald (2019).

All five pectinariid genera occur in European waters, and are represented by seven species and one subspecies (Gil 2011). Among these, *Amphictene auricoma mediterranea* Nilsson, 1928 and *Lagis neapolitana* (Claparède, 1869) have only been reported from the Mediterranean Sea. The remaining six species, namely *Amphictene auricoma* (O.F. Müller, 1776), *Cistenides granulata* (Linnaeus, 1767), *C. hyperborea* Malmgren, 1865, *Lagis koreni* Malmgren, 1866, *Pectinaria belgica* (Pallas, 1766) and *Petta pusilla* Malmgren, 1866, are distributed across the NE Atlantic coasts; *A. auricoma*, *C. granulata*, *C. hyperborea* and *L. koreni* have been also recorded in Iceland prior to the present study (Wesenberg-Lund 1951).

Among other characters, the terebelliform polychaetes, and the Pectinariidae in particular, are characterized by having neurochaetae modified as uncini for anchoring the body within the tube, in order to avoid the risk of predation (Hutchings 2000; Rouse & Pleijel 2001; Garraffoni & Camargo 2006). Nogueira *et al.* (2019) (based on Rouse & Pleijel 2001; Hutchings & Peart 2002; Nogueira *et al.* 2010; Zhang & Qiu 2017) refer to two types of neurochaetal uncini in Pectinariidae: A) *Pectinate uncini*, with an anterior peg (hereafter ‘subrostral process’) composed of densely packed denticles, uncinal teeth (hereafter ‘main teeth’) long and thin, arranged in longitudinal rows, and main fang (hereafter ‘rostrum’) absent; and B) *Avicular uncini*, with rostrum present and secondary teeth in transverse rows and progressively shorter distally (hereafter ‘capitium’).

In this paper, we study the diversity and distribution of the Pectinariidae around Iceland after a number of samples collected during the BIOICE cruises. The aforementioned species are reported here again for Iceland and remarks about their taxonomy and ecology are provided. Furthermore, we have also studied the uncini in specimens of different body sizes for all four species; this was done by using the SEM in order to test 1) whether there were morphological differences supposedly related to ontogeny and 2) whether this uncini variability showed any pattern both within a species and among species and therefore could be used as a taxonomic character.

Material and methods

This study is based on material collected in the framework of the BIOICE project. A total of 1066 pectinariid specimens were found in 43 samples; some specimens in eight samples could not be identified because of their small size or bad preservation (e.g., fragmented body). Samples were initially fixed in 10% formalin buffered with borax, and then preserved in 70% ethanol. Specimens were picked from

samples by the staff of the Sandgerdi Marine Centre (SMC, Iceland). Most of the material examined was deposited in the collections of the Icelandic Institute of Natural History (IINH, Reykjavík); several selected specimens were deposited in the Museo Nacional de Ciencias Naturales (MNCN, Madrid).

Abiotic data of all BIOICE samples are available from the following link: http://utgafa.ni.is/greinar/BIOICE_station_list_91-04_Paper_A2.pdf

Identification of specimens and stereomicrographs were done using an Olympus SZX9 stereo microscope at the Facultade de Ciencias, Universidade da Coruña (UDC, Spain). Specimens used for examination with SEM were prepared by critical point drying, covered with gold in a BAL-TEC SCD 004 evaporator and examined and photographed under a JEOL JSM-6400 scanning electron microscope (SEM) at the Servizos de Apoio á Investigación (SAI–UDC, Spain).

Nomenclature for morphological characters, genera, species and synonyms mostly follows Holthe (1986) and Hutchings & Peart (2002). Each species is related to water mass(es) as described by Brix & Svavarsson (2010) according to geographic location (coordinates), depth and temperature.

List of morphological abbreviations

al	=	anal lobe
alcf	=	anal lobe ciliary field
ap	=	anal papilla
bp	=	basal prow
br	=	branchia
bt	=	buccal tentacle
cap	=	capitium
cp	=	cuticular pores
cv	=	cephalic veil
cvc	=	cephalic veil cirri
dc	=	dorsal chaetae
ga	=	glandular area
mt	=	main tooth/teeth
or	=	opercular rim
pa	=	paleae
ro	=	rostrum
sbcf	=	scaphal basis ciliary field
scb	=	scaphal border
sch	=	scaphal hooks
scl	=	scaphal lobes
srp	=	subrostral process
tc	=	tentacular cirrus
un	=	unciniger
unXd	=	un, unciniger; X, number of uncinigerous segment; d, dorsalmost uncini within row
unXm	=	un, unciniger; X, number of uncinigerous segment; m, median uncini within row
unXv	=	un, unciniger; X, number of uncinigerous segment; v, ventralmost uncini within row
vu	=	ventral uncini

List of water mass abbreviations

CW	=	Coastal Waters
MEIW	=	Modified East Icelandic Water
MNAW	=	Modified North Atlantic Waters
NSAIW	=	Norwegian Sea Arctic Intermediate Waters

Results

Taxonomy

The study of the samples collected during the BIOICE project yielded four pectinariid species belonging to three genera. The most abundant species was *Lagis koreni* (801 specs; 75.1% of the total pectinariid specimens identified), followed by *Amphictene auricoma* (117 specs; 11.0%), *Cistenides hyperborea* (125 specs; 11.7%) and *C. granulata* (23 specs; 2.2%). All four species show a well-defined geographical distribution around Iceland corresponding to narrow ranges of temperature and depth, but being mostly present in temperate waters at depths of between 100 and 300 m (Figs 1–2); there were some exceptions to this pattern corresponding to samples with *C. hyperborea* and *L. koreni*. Abundance and distribution around Iceland of each species and taxonomic remarks are presented and discussed below.

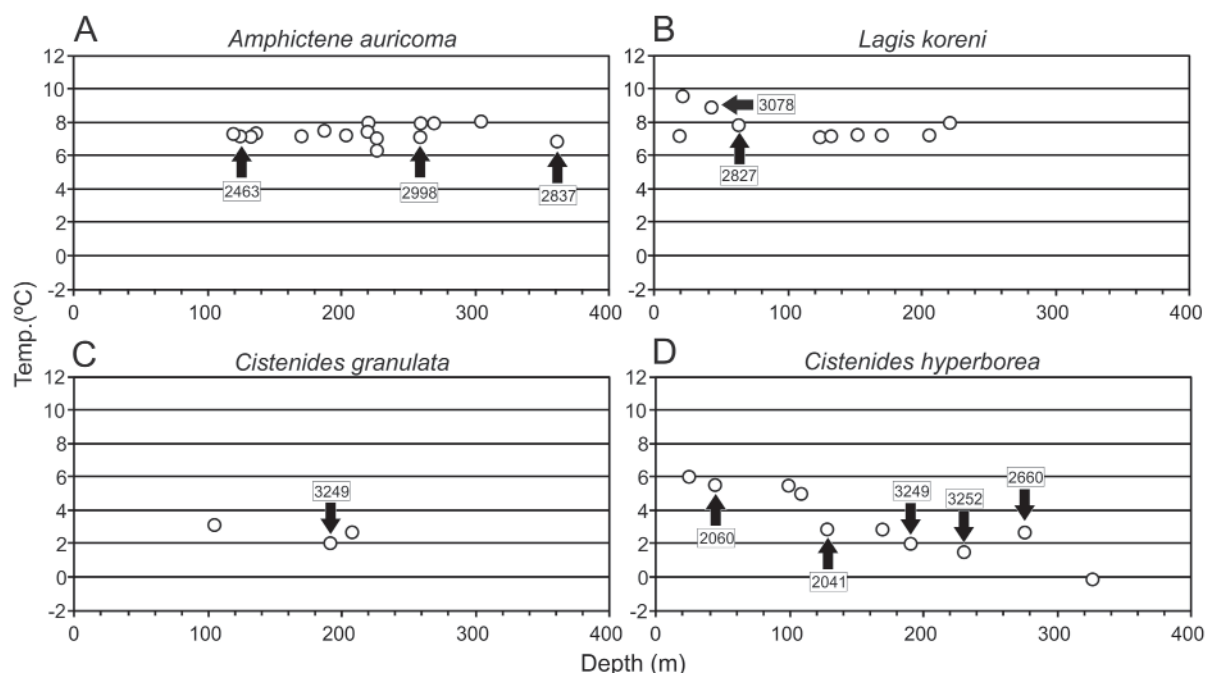


Fig. 2. Temperature vs depth data of the BIOICE samples where specimens of Pectinariidae were found. **A.** *Amphictene auricoma* (O.F. Müller, 1776). **B.** *Lagis koreni* Malmgren, 1866. **C.** *Cistenides granulata* (Linnaeus, 1767). **D.** *Cistenides hyperborea* Malmgren, 1866. Specimens studied under SEM are indicated with black arrows and sample numbers.

Phylum Annelida Lamarck, 1809
Family Pectinariidae Quatrefages, 1866

Genus *Amphictene* Lamarck, 1818

Type species and type locality

Amphictene auricoma Müller, 1776. Denmark; type probably lost or never designated (Holthe 1986).

Remarks

The genus is characterized according to Hutchings & Peart (2002) by having neurochaetal uncini with major teeth arranged in two rows. However, an examination of BIOICE specimens of *A. auricoma* suggests that this character could be reassessed (see Discussion below).

Amphictene auricoma (O.F. Müller, 1776)

Figs 1A, 2A, 3A, 4A–B, 5–7

Amphitrite auricoma O.F. Müller, 1776: 216.

Amphictene auricoma – Malmgren 1865: 357. — Hutchings & Peart 2002: 103, table 1.

Pectinaria auricoma – Hesse 1917: 78. — Jirkov & Leontovich 2013: 220, key.

Pectinaria (Amphictene) auricoma – Fauvel 1927: 222. — Holthe 1986: 22. — Hartmann-Schröder 1996: 480. — Kirkegaard 1996: 280.

Material examined (117 specimens, 11.0% of all specimens identified, in 18 samples)

ICELAND – **North Western Fjords** • 3 specs; BIOICE station 489, sample 2946; 65°47'90" N, 25°38'70" W; 28 Aug. 1996; 6.20°C; 35.05 ppm; 228 m depth; sandy silt; IINH-40181 • 1 spec.; BIOICE station 489, sample 2947; 65°47'91" N, 25°38'68" W; 29 Aug. 1996; 6.20°C; 35.05 ppm; 227 m depth; sediment unknown; IINH-40211. – **South West to South East coast** • 1 spec.; BIOICE station 968, sample 2209; 63°59'01" N, 23°34'13" W; 3 Sep. 1992; 7.29°C; 35.07 ppm; 137 m depth; fine mud; IINH-40117 • 3 specs; BIOICE station 553, sample 2388; 63°30'10" N, 22°03'70" W; 30 Jun. 1993; 7.13°C; 35.10 ppm; 171 m depth; silty sand; IINH-27841 • 1 spec.; BIOICE station 564, sample 2417; 63°09'90" N, 21°11'80" W; 2 Jul. 1993; 7.08°C; 35.11 ppm; 259 m depth; sandy silt; IINH-40113 • 5 specs; BIOICE station 572, sample 2440; 63°20'21" N, 19°49'55" W; 4 Jul. 1993; 6.87°C; 35.10 ppm; 228 m depth; sandy silt; MNCN 16.01/17982 • 1 spec.; BIOICE station 572, sample 2441; 63°20'17" N, 19°49'60" W; 4 Jul. 1993; 6.87°C; 35.10 ppm; 228 m depth; sandy silt; IINH-40119 • 1 spec.; BIOICE station 582, sample 2460; 63°29'65" N, 21°39'10" W; 5 Jul. 1993; 7.10°C; 35.08 ppm; 125 m depth; sediment unknown; IINH-40127 • 42 specs; BIOICE station 583, sample 2463; 63°25'40" N, 21°39'89" W; 5 Jul. 1993; 7.12°C; 35.08 ppm; 133 m depth; silty sand and gravel; IINH-40155 • 4 specs;

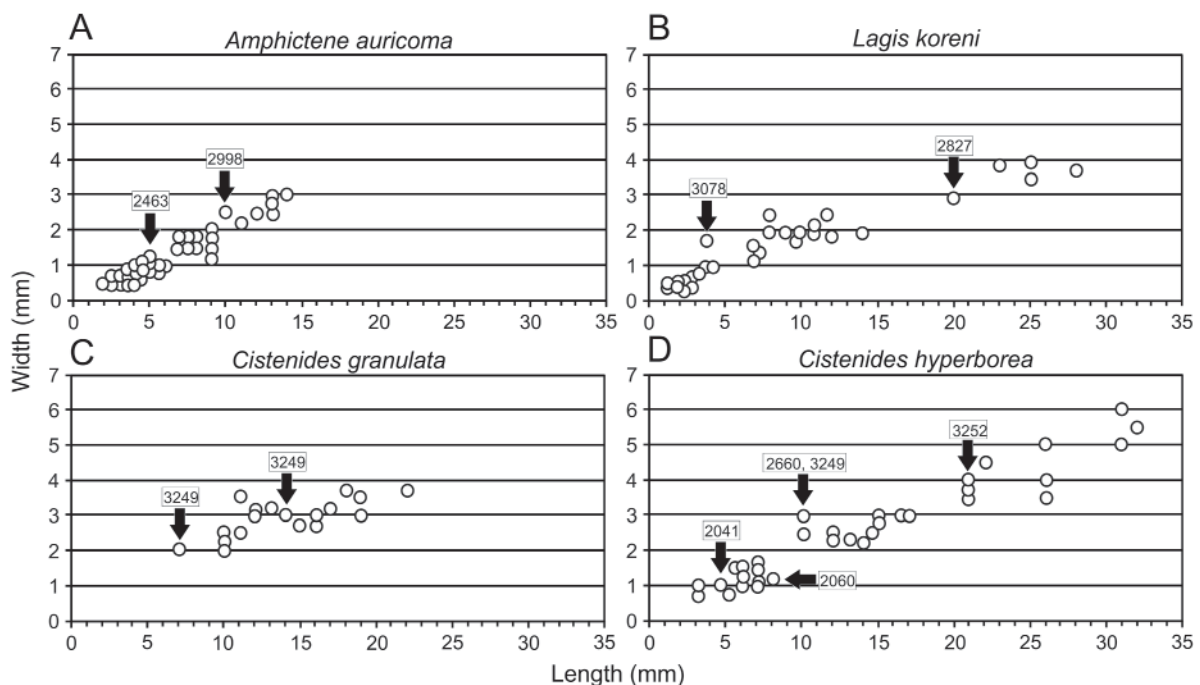


Fig. 3. Biometric features of the Pectinariidae specimens in BIOICE samples. **A.** *Amphictene auricoma* (O.F. Müller, 1776). **B.** *Lagis koreni* Malmgren, 1866. **C.** *Cistenides granulata* (Linnaeus, 1767). **D.** *Cistenides hyperborea* Malmgren, 1866. Specimens studied under SEM are indicated with black arrows and sample numbers.

BIOICE station 715, sample 2817; 63°14'75" N, 17°50'62" W; 24 Aug. 1995; 7.18°C; 35.14 ppm; 204 m depth; sediment unknown; IINH-40165 • 5 specs; BIOICE station 717, sample 2824; 63°30'12" N, 17°42'07" W; 25 Aug. 1995; 7.24°C; 35.13 ppm; 120 m depth; gravelly sand; IINH-40166 • 6 specs; BIOICE station 723, sample 2837; 63°16'66" N, 16°53'52" W; 26 Aug. 1995; 6.74°C; 35.12 ppm; 601 m depth; sandy gravel and corals; IINH-40170 • 18 specs; BIOICE station 275, sample 2998; 63°38'20" N, 14°43'50" W; 5 Jul. 1997; 7.76°C; 35.19 ppm; 264 m depth; silty sand; IINH-40213 • 5 specs; BIOICE station 275, sample 2999; 63°38'20" N, 14°43'60" W; 5 Jul. 1997; 7.76°C; 35.19 ppm; 269 m depth; sediment unknown; IINH-40293 • 14 specs; BIOICE station 299, sample 3061; 63°59'49" N, 14°09'21" W; 10 Jul. 1997; 7.59°C; 35.16 ppm; 221 m depth; sediment unknown; IINH-40461 • 4 specs; BIOICE station 725, sample 3257; 63°20'10" N, 19°52'20" W; 11 Sep. 2001; 7.92°C; 35.19 ppm; 221 m depth; sediment unknown; IINH-40462 • 1 spec.; BIOICE station 734, sample 3275; 63°23'10" N, 16°16'20" W; 15 Sep. 2001; 7.97°C; 35.20 ppm; 305 m depth; sediment unknown; IINH-40463 • 1 spec.; BIOICE station 407, sample 3610; 63°58'80" N, 25°30'48" W; 12 Sep. 2003; 7.43°C; 35.17 ppm; 188 m depth; silty sand; IINH-40464.

Occurrence

From off West and South of Reykjanes Peninsula to Höfn at SE; two additional samples from off north-western fjords (Fig. 1). Depth range: 120 to 305 m (sample 2837: 601 m, not shown); bottom temperature range: 6.20 to 7.97°C (Fig. 2A). Water mass/es: MNAW.

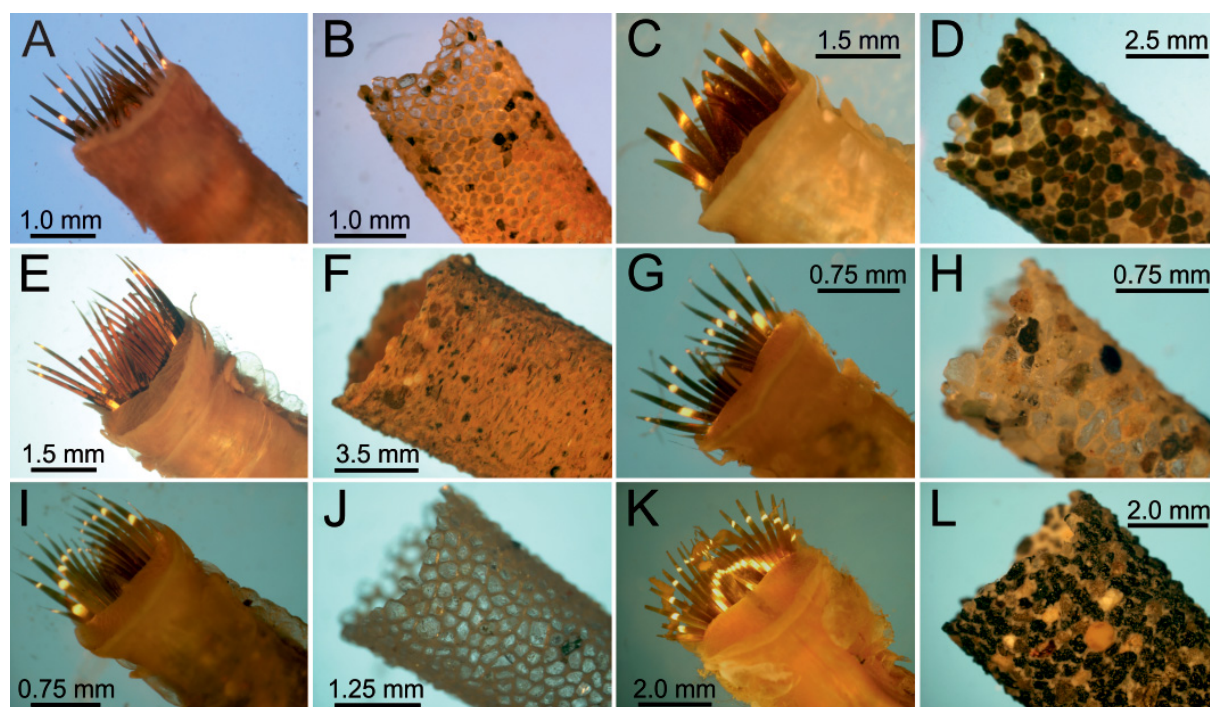


Fig. 4. Stereomicrographs of anterior body ends (dorsal view) and anterior end of the tube. **A–B.** *Amphictene auricoma* (O.F. Müller, 1776) (BIOICE sample 2946, IINH-40181). **C–D.** *Cistenides granulata* (Linnaeus, 1767) (sample 2126, IINH-40466). **E–H.** *Cistenides hyperborea* Malmgren, 1866 (E–F: sample 3252, IINH-40477; G–H: sample 2060, IINH-40471). **I–L.** *Lagis koreni* Malmgren, 1866 (I–J: sample 2557; K–L: sample 2827, IINH-40485).

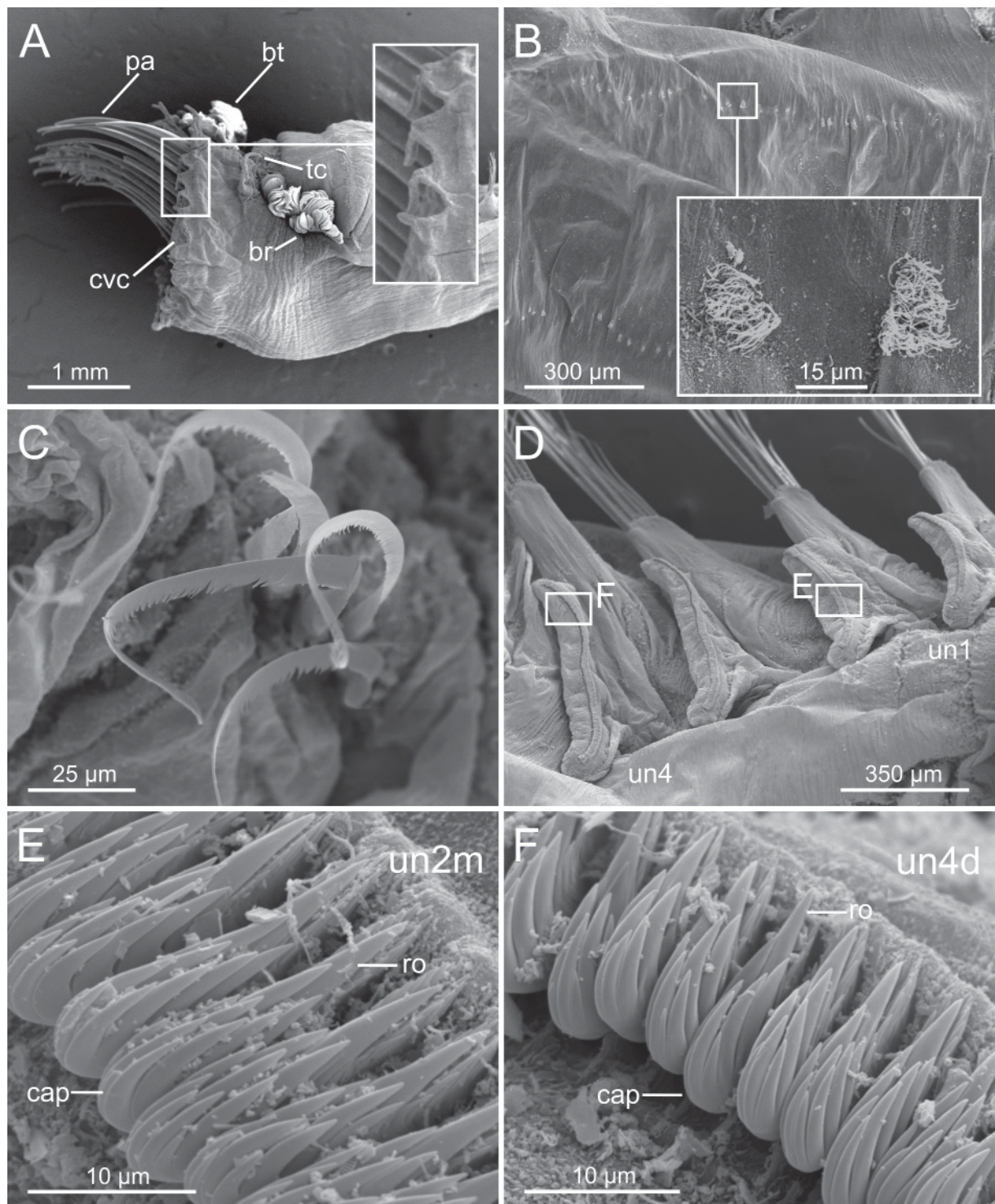


Fig. 5. *Amphictene auricoma* (O.F. Müller, 1776). SEM micrographs from two large specimens (BIOICE sample 2998, IINH-40213). **A.** Anterior end, left lateral view (framed: cirri of opercular rim, detail). **B.** Ventral rows of ciliated tufts (framed: two tufts, detail). **C.** Notochaetae; distal end. **D.** Uncinigers 1 to 4. **E.** Unciniger 2, median uncini (framed in D). **F.** Unciniger 4, dorsal uncini (framed in D).

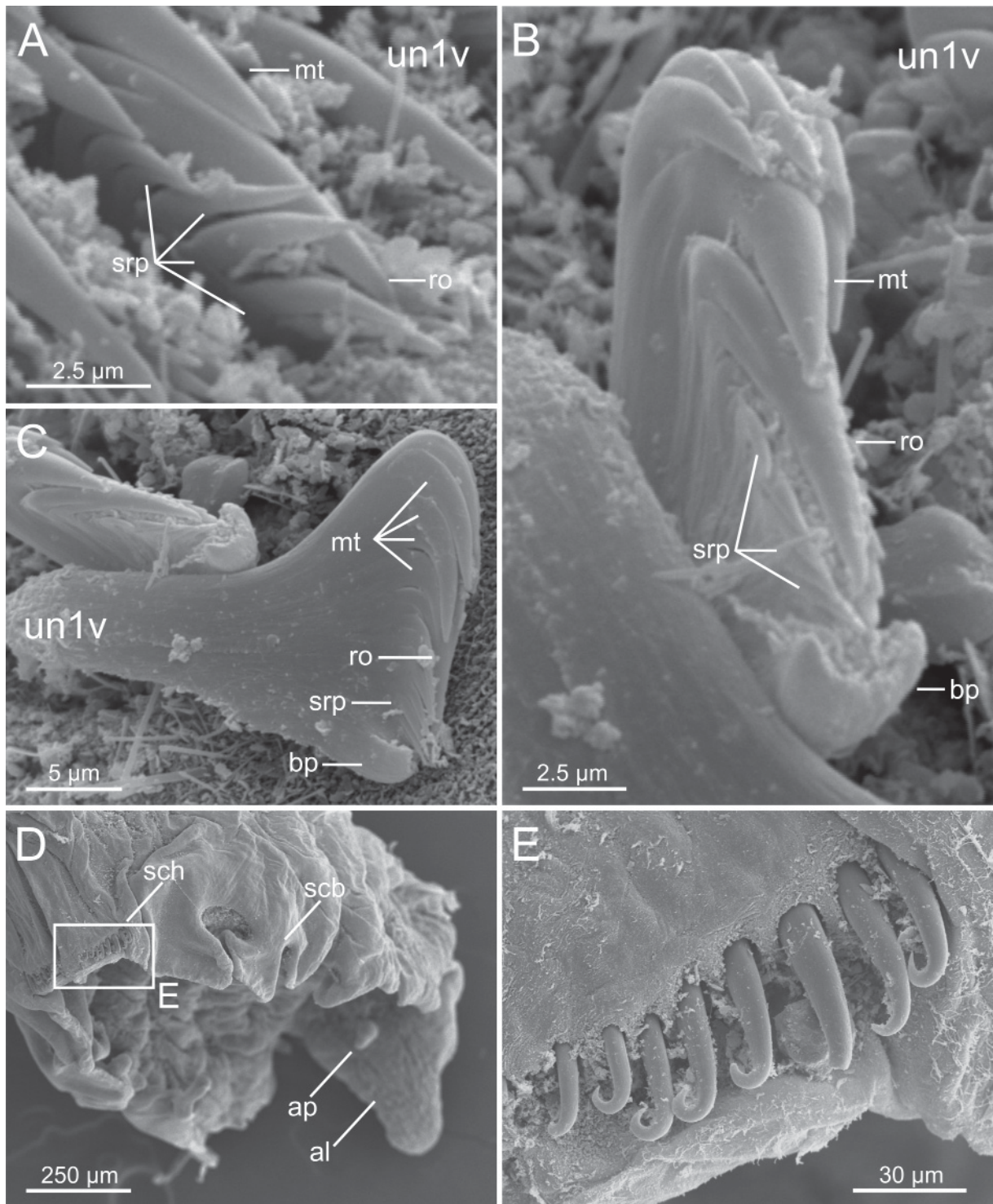


Fig. 6. *Amphictene auricoma* (O.F. Müller, 1776). SEM micrographs from a large specimen (BIOICE sample 2998, IINH-40213). **A–C.** Unciniger 1, ventral uncinus, frontolateral, frontal and lateral views, respectively. **D.** Scaphe. **E.** Scaphal hooks (framed in D).

Remarks

Species of *Amphictene* bear a characteristic crenulated opercular rim, that is not present in other genera (Figs 4A, 5A). *Amphictene auricoma* – type locality: Denmark (Holthe 1986) – has been the only species of the genus reported in North Atlantic waters and also bears thick paleae with sharp tips (Fig. 5A), scaphe with crenulated margins (Fig. 6D), 17 uniramous and 13 biramous chaetigers, and distally curved scaphal hooks (Fig. 6E). However, Hutchings & Peart (2002: table 1) pointed out that there is “a considerable variation recorded for this species” in the North Atlantic. General morphology of BIOICE specimens agree well with previous descriptions and the aforementioned characters; however, SEM examination revealed that several pygidial and uncini features differ from what was described for *A. auricoma*. First, Holthe (1986: fig. 4c) and Hartmann-Schröder (1996: fig. 234c) mentioned a long “anal tongue” and “anal cirrus”, corresponding to the “anal lobe” and “anal papilla” (after Hutchings & Peart 2002), that are much shorter in BIOICE specimens (Fig. 6D).

On the other hand, the uncini of *A. auricoma* have been described as having 1–2 vertical rows of main teeth (cf. Hartmann-Schröder 1971: fig. 157b, and later works: Holthe 1986: fig. 4e; Hartman-Schröder 1996: fig. 234b). However, SEM micrographs of BIOICE material shows a well-defined rostrum, surrounded in all its length by long teeth forming, in turn, a subrostral process (Fig. 6A–B); the rostrum is surmounted by a capitium constituted by many large teeth not arranged in vertical lines (as described in the original description) but with a typical avicular arrangement (Figs 5E–F, 6A–B). Anyway, these uncini may resemble in lateral view (Fig. 6C) the typical ones of *A. auricoma* and this may have led previous authors to confusion.

In addition, uncinal denticulation shows dorsoventral variation within chaetigers depending on body size. Thus, in large specimens (Fig. 3A) uncini seem avicular along the whole torus (Fig. 5E–F); in smaller ones, the avicular uncini are only found in the dorsal part of the torus (Fig. 7A, C) while in the ventral side all capitium teeth are similar in size and covering most of the rostrum (only showing the pointed distal end; Fig. 7B, D). This pattern is even more evident in the last unciniger, where the capitium teeth seem mixed with those of the subrostral process and leaving a small opening through which the tip of the rostrum is visible (Fig. 7E).

Finally, we have found two longitudinal rows of ciliated tufts/patches along the whole ventral body surface (Fig. 5B) that have not been mentioned in the literature to the best of our knowledge, and those tufts may be related to water irrigation within the tube.

Genus *Cistenides* Malmgren, 1865

Type species and type locality

Sabella granulata Linnaeus, 1767. Northern Europe; type never designated (Holthe 1986).

Remarks

The diagnosis of the genus in Hutchings & Peart (2002) includes “... chaetigers 4 to 16 biramous with notopodia, neuropodia, noto chaetae and neurochaetae, chaetiger 17 with notopodia and noto chaetae only (17/13) ...” and the presence of a single row of teeth on the uncini as well. Considering the last character, Hutchings & Peart (2002) transferred four species previously referred to *Cistenides*, namely *C. aegyptia* Savigny, 1822, *C. chilensis* (Nilsson, 1928), *C. gouldii* Verrill, 1874 and *C. regalis* Verrill, 1900, to the genus *Pectinaria*, because of having uncini with two rows of teeth, a character supposedly typical of this genus. We think that both characters could be reconsidered (see Discussion below).

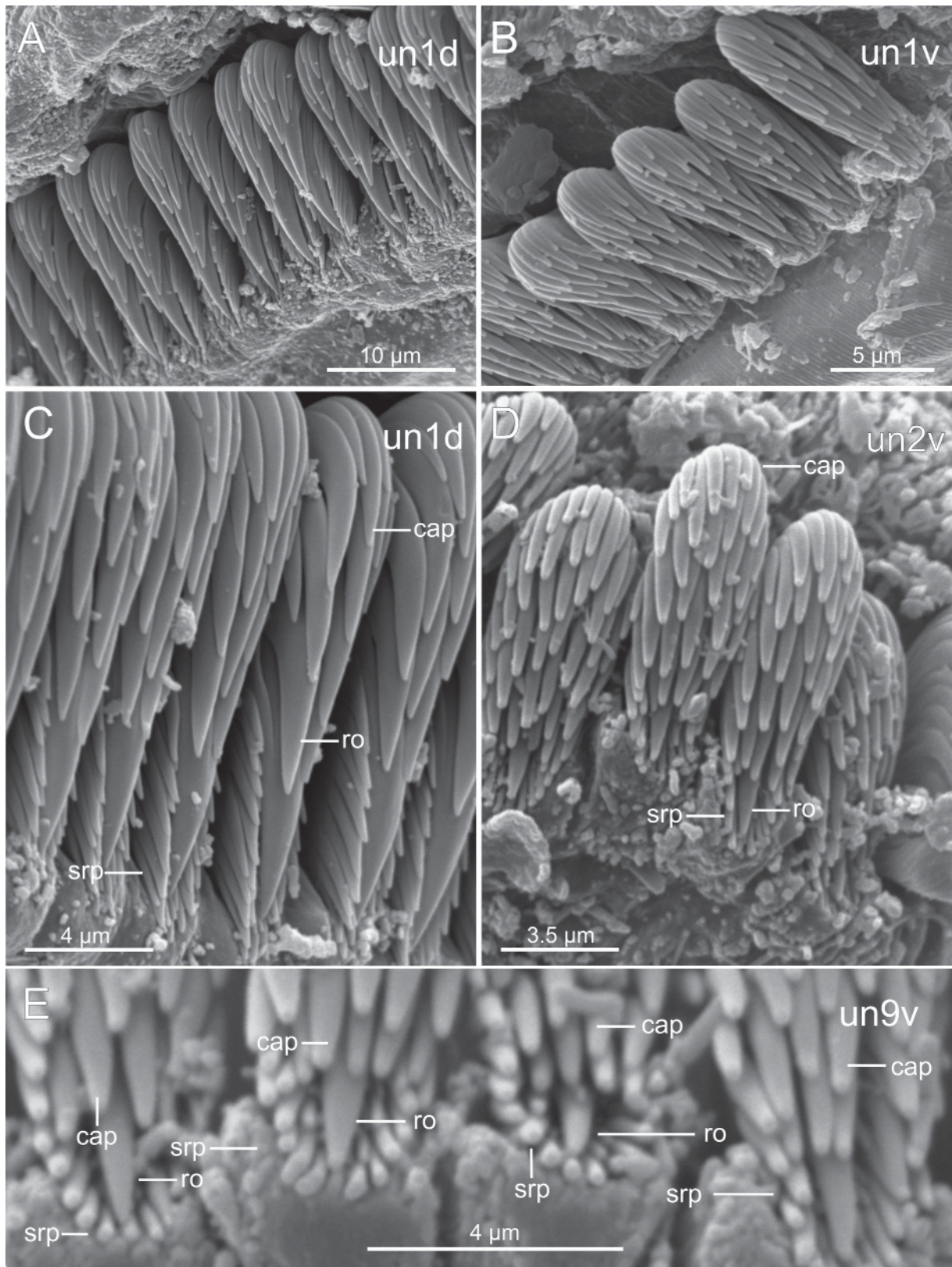


Fig. 7. *Amphictene auricoma* (O.F. Müller, 1776). SEM micrographs from a small specimen (BIOICE sample 2463, IINH-40481). **A–B.** Unciniger 1, dorsal and ventral uncini, respectively. **C–D.** The same in closer view. **E.** Unciniger 9, ventral uncini, fronto-inferior area.

Cistenides granulata (Linnaeus, 1767)

Figs 1C, 2C, 3C, 4C–D, 8–9

Sabella granulata Linnaeus, 1767: 1268.

Pectinaria granulata – Hessle 1917: 77.

Pectinaria (Cistenides) granulata – Nilsson 1928: 28. — Pettibone 1954: 312, fig. 35i–k. — Holthe 1986: 24. — Hartmann-Schröder 1996: 482.

Material examined (23 specimens, 2.2% of total specimens identified, in three samples)

ICELAND – **North and North East coast** • 7 specs; BIOICE station 30, sample 2042; 65°49'56" N, 14°32'94" W; 24 Jul. 1991; 3.1°C; salinity unknown; 105 m depth; sediment unknown; MNCN 16.01/17988 • 9 specs; BIOICE station 27, sample 2126; 66°59'51" N, 18°49'82" W; 8 Jul. 1992; 2.7°C; 34.86 ppm; 208 m depth; sandy gravel and stones; IINH-40466 • 7 specs; BIOICE station 23, sample 3249; 65°50'34" N, 12°01'27" W; 14 Jul. 2001; 1.92°C; 34.87 ppm; 192 m depth; sediment unknown; IINH-40467.

Occurrence

From off northern and north-eastern coast of Iceland. Depth range: 105 to 208 m (Fig. 1C); bottom temperature range: 1.92 to 3.10°C (Fig. 2C). Water mass/es: MEIW.

Remarks

All diagnostic characters for this species – type locality: Northern Europe (Holthe 1986) – as stated by Holthe (1986) are present in BIOICE material: shape and number of paleae and cephalic veil (Figs 4C, 8A, 9A), scaphal shape, anal lobe and anal papilla (Fig. 8F). Ventral uncini agree well with those present in the genus, not showing variations within uncinigers nor along the body both in large (Fig. 8B–E) and small (Fig. 9B–F) specimens; there is only one row of teeth becoming progressively smaller in size from the uncinus base (at the level of the subrostral process) towards the distal end. This lack of variation, which has been considered the typical pattern in the family, was only found in *C. granulata* among the four pectinariid species in BIOICE samples.

Cistenides hyperborea Malmgren, 1865

Figs 1D, 2D, 3D, 4E–H, 10–15

Cistenides hyperborea Malmgren, 1865: 360.

Pectinaria hyperborea – Hessle 1917: 76.

Pectinaria (Cistenides) hyperborea – Nilsson 1928: 31. — Pettibone 1954: 312, fig. 35c–h. — Holthe 1986: 25. — Hartmann-Schröder 1996: 482.

Material examined (125 specimens, 11.7% of total specimens identified, in ten samples)

ICELAND – **North coast** • 1 spec.; BIOICE station 1, sample 2001; 65°21'19" N, 13°47'61" W; 19 Jul. 1991; 6.0°C; salinity unknown; 26 m depth; organic silt; IINH-40468 • 3 specs; BIOICE station 3, sample 2005; 65°20'88" N, 13°18'64" W; 19 Jul. 1991; 2.8°C; salinity unknown; 171 m depth; sandy silt and stones; MNCN 16.01/18015 • 1 spec.; BIOICE station 29, sample 2041; 65°48'61" N, 14°34'01" W; 24 Jul. 1991; 2.8°C; salinity unknown; 129 m depth; sediment unknown; IINH-40470 • 54 specs; BIOICE station 2, sample 2060; 66°00'61" N, 17°31'78" W; 2 Jul. 1992; temperature unknown; salinity unknown; 48 m depth; soft sediment; IINH-40471 • 3 specs; BIOICE station 3, sample 2064; 66°02'33" N, 17°32'21" W; 2 Jul. 1992; 5.4°C; 34.69 ppm; 102 m depth; mud; IINH-40472 • 16 specs;

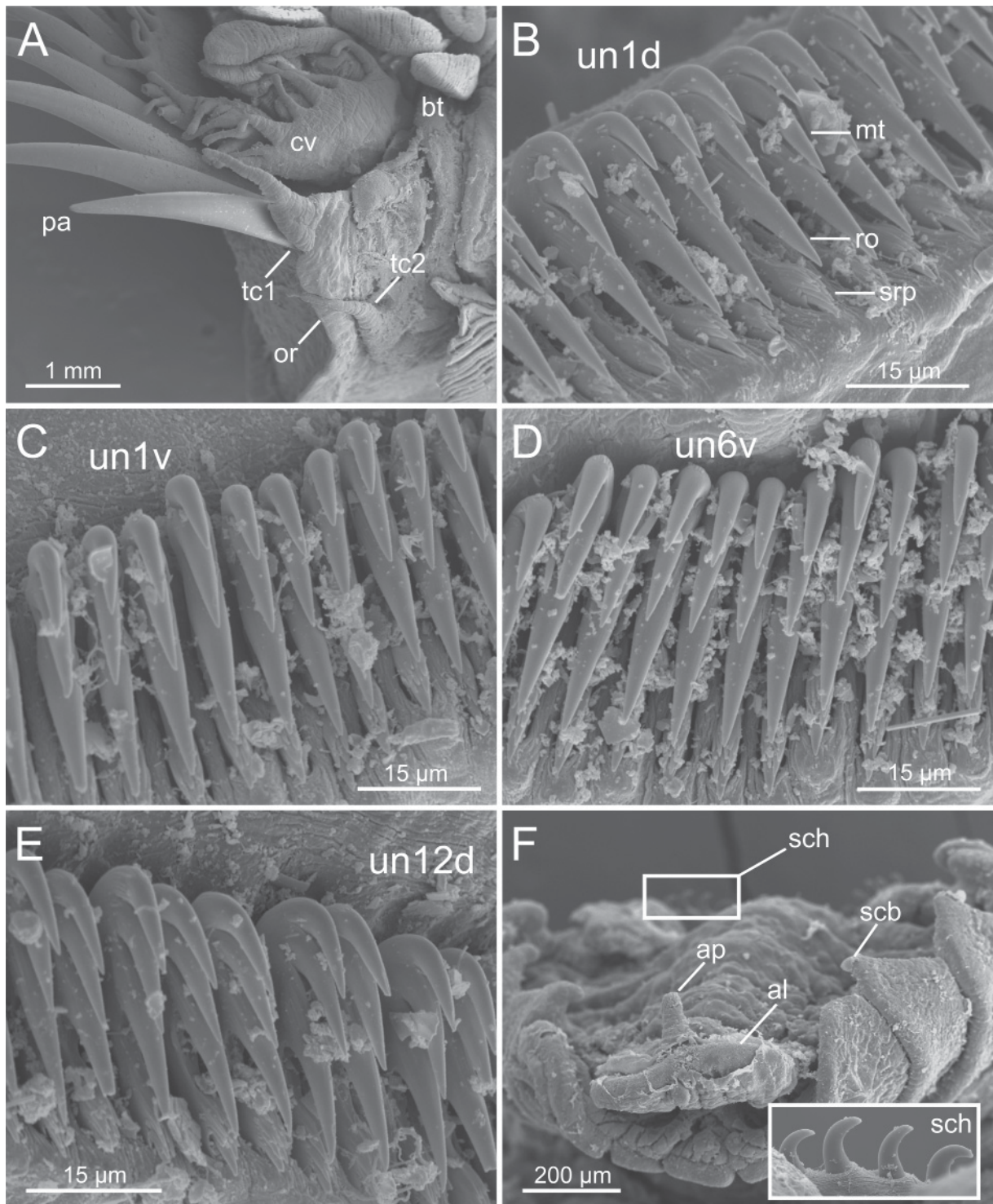


Fig. 8. *Cistenides granulata* (Linnaeus, 1767). SEM micrographs from two large specimens (BIOICE sample 3249, IINH-40467). **A.** Anterior end, right lateral view. **B–C.** Unciniger 1, dorsal and ventral uncini, respectively. **D.** Unciniger 6, ventral uncini. **E.** Unciniger 12, dorsal uncini. **F.** Scaphe, postero-dorsal view (framed: scaphal hooks, detail).

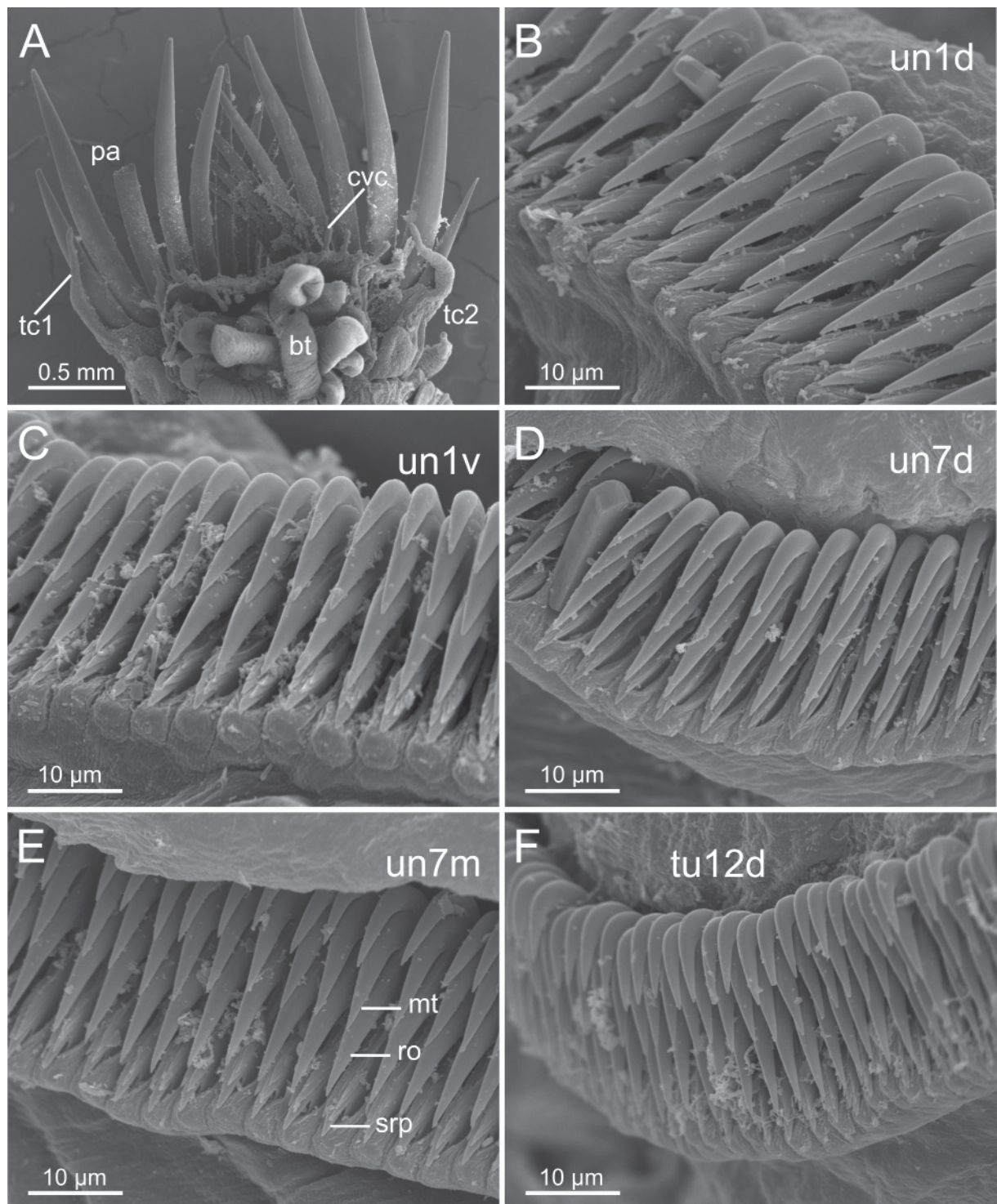


Fig. 9. *Cistenides granulata* (Linnaeus, 1767). SEM micrographs from a small specimen (BIOICE sample 3249, IINH-40467). **A.** Anterior end, ventral view. **B–C.** Unciniger 1, dorsal and ventral uncini, respectively. **D–E.** Unciniger 7, dorsal and median uncini, respectively. **F.** Unciniger 12, dorsal uncini.

BIOICE station 16, sample 2097; 66°36'92" N, 18°14'42" W; 5 Jul. 1992; 4.88°C; 34.89 ppm; 110 m depth; sand and stones; IINH-40473 • 17 specs; BIOICE station 17, sample 2660; 67°14'41" N, 15°28'42" W; 15 Jul. 1994; 2.69°C; 34.90 ppm; 277 m depth; sandy silt; IINH-40474 • 5 specs; BIOICE station 5, sample 3108; 67°45'48" N, 18°30'85" W; 21 Aug. 1999; -0.21°C; 34.88 ppm; 328 m depth; sediment unknown; IINH-40475 • 23 specs; BIOICE station 23, sample 3249; 65°50'34" N, 12°01'27" W; 14 Jul. 2001; 1.92°C; 34.87 ppm; 192 m depth; sediment unknown; IINH-40476 • 24 specs; BIOICE station 24, sample 3252; 65°45'94" N, 12°16'39" W; 14 Jul. 2001; 1.48°C; 34.86 ppm; 232 m depth; sediment unknown; IINH-40477.

Occurrence

From off northern and north-eastern coast of Iceland (Fig. 1D). Depth range: 26 to 328 m; bottom temperature range: -0.21 to 6.00°C (Fig. 2D). Water mass/es: NSAIW, CW and MNAW.

Remarks

Cistenides hyperborea – type locality: Greenland and Spitzbergen (Holthe 1986) – is a poorly known species; the original description does not indicate the number of rows of teeth present on the uncini, although 1 to 3 rows have previously been mentioned for specimens assigned to this taxon (Hutchings & Peart 2002). Because type material was never designated (Holthe 1986), Hutchings & Peart (2002) suggested that specimens from the type locality need to be re-examined to ascertain the generic identity of this species.

The examination of the Icelandic specimens here identified as *C. hyperborea*, covering a wide range of body sizes, shows a considerable variation in the shape of the neuropodial uncini, contrary to the expected, according to the description of the species. BIOICE specimens agree with previous descriptions (e.g., Pettibone 1954; Holthe 1986; Hutchings & Peart 2002) in: 1) having a smooth dorsal brim and cephalic veil free from operculum, provided with several well-defined cirri (Fig. 10A); 2) the relation between chaetigers with notochaetae *versus* those with also ventral uncini is 17/12; 3) the paleae are long, numerous, straight and with pointed tips (Figs 10A, 12A); 4) the scaphe has small lateral lobes (Fig. 11A) with a short and rounded anal lobe and anal papilla (Fig. 11B), and scaphal hooks are distally bent (Fig. 11C–E). On the other hand, the aforementioned characters show, however, differences related to the size of the specimen. For instance, medium-sized and small individuals show a strait scaphe border with a verrucose anal lobe (Fig. 11E–F), thinner paleae (Figs 12A, 13A) and the tip of the scaphal hooks is strongly curved (compare Figs 12F, 13F, 14F).

The number and arrangement of teeth in the neuropodial uncini shows differences among BIOICE individuals, contrary to that observed in *C. granulata*. Two different types of uncini can be distinguished:

- 1) Uncini with 2–3 main teeth arranged in a vertical row. This is the typical teeth arrangement in the *pectinate* type for this species and the genus as well. We found this pattern in large specimens (Figs 3D, 10).
- 2) Uncini provided with a large basal tooth (rostrum) and an upper group of short teeth (capitium). This agrees with the typical *avicular* type present in many species of Terebellomorpha and found here in smaller specimens (Figs 3D, 12–13).

SEM examination of BIOICE specimens seems to support the hypothesis that these differences are related to body size even though we did not examine the whole range of sizes (Fig. 3D). Uncini of large specimens (sample 3252, Fig. 10) do not show a variation either within an individual torus or along the body; on the contrary, mid-sized specimens (sample 2660, Fig. 12) show a progressive increase in the number of upper teeth, mostly in ventral uncini (Fig. 12C, E) and especially in posterior chaetigers (Fig. 12E), the latter having a well-defined capitium; dorsal uncini show, in turn, the usual shape as

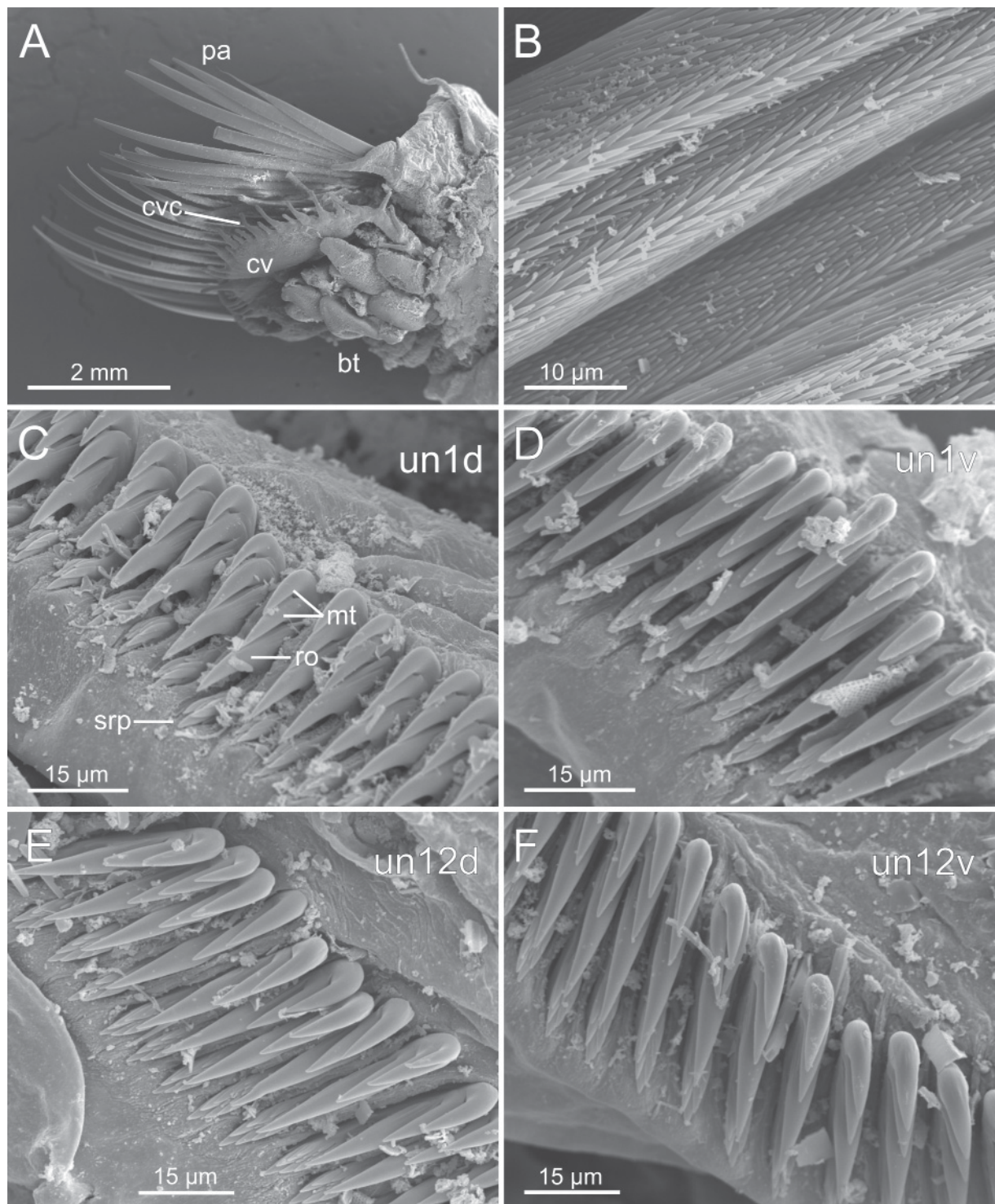


Fig. 10. *Cistenides hyperborea* Malmgren, 1866. SEM micrographs from three large specimens (BIOICE sample 3252, IINH-40477). **A.** Anterior end, left ventrolateral view. **B.** Notochaetal scale covering, detail. **C–D.** Unciniger 1, dorsal and ventral uncini, respectively. **E–F.** Unciniger 12, dorsal and ventral uncini, respectively.

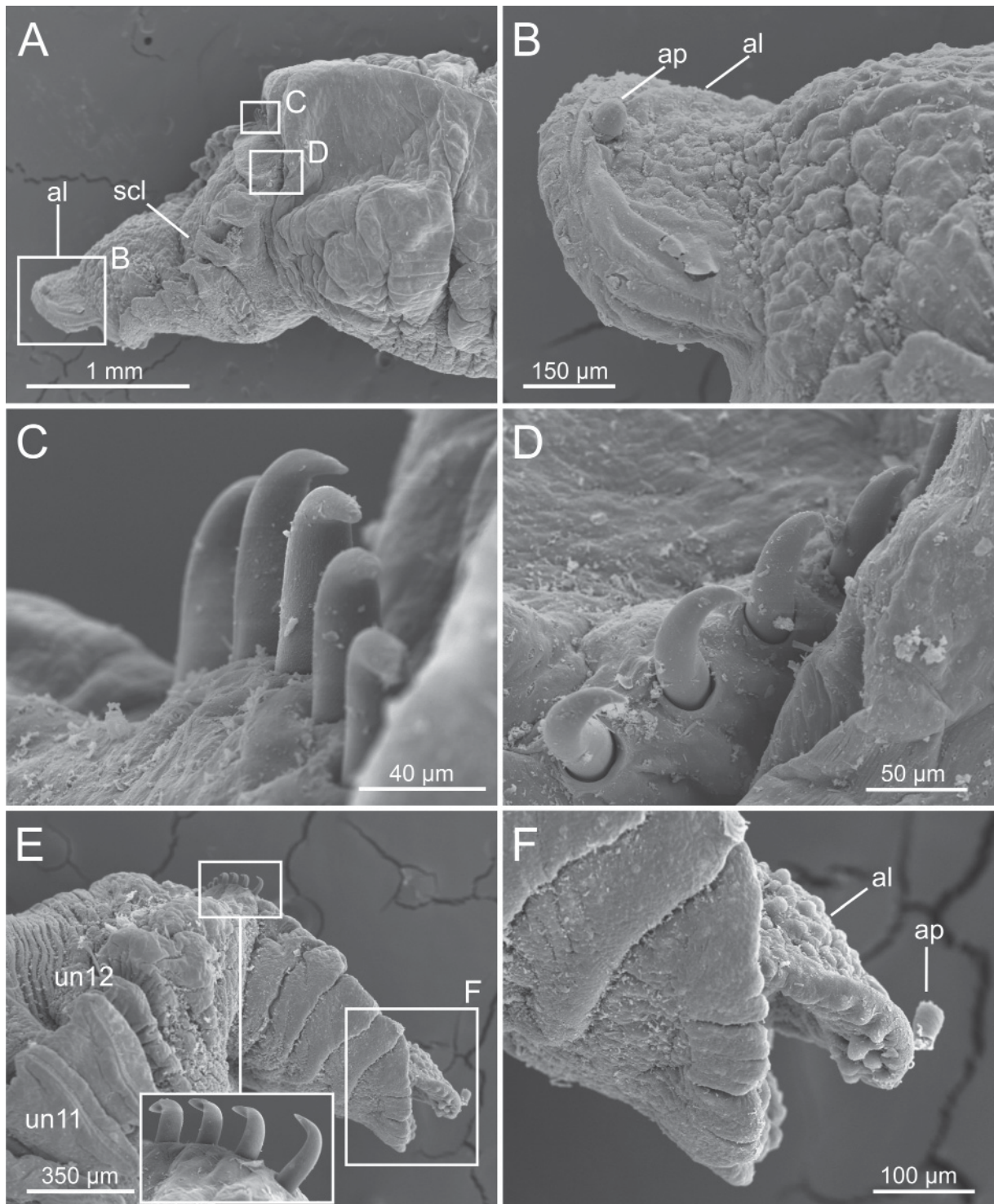


Fig. 11. *Cistenides hyperborea* Malmgren, 1866. SEM micrographs from four large specimens (A–D: BIOICE sample 3252, IINH-40477; E–F: sample 3249, IINH-40476). **A.** Scaphe, left lateral view. **B.** Anal lobe and anal papilla (framed in A). **C–D.** Scaphal hooks (framed in A). **E.** Posterior end (framed: scaphal hooks, detail). **F.** Scaphe and anal lobe and papilla (framed in E).

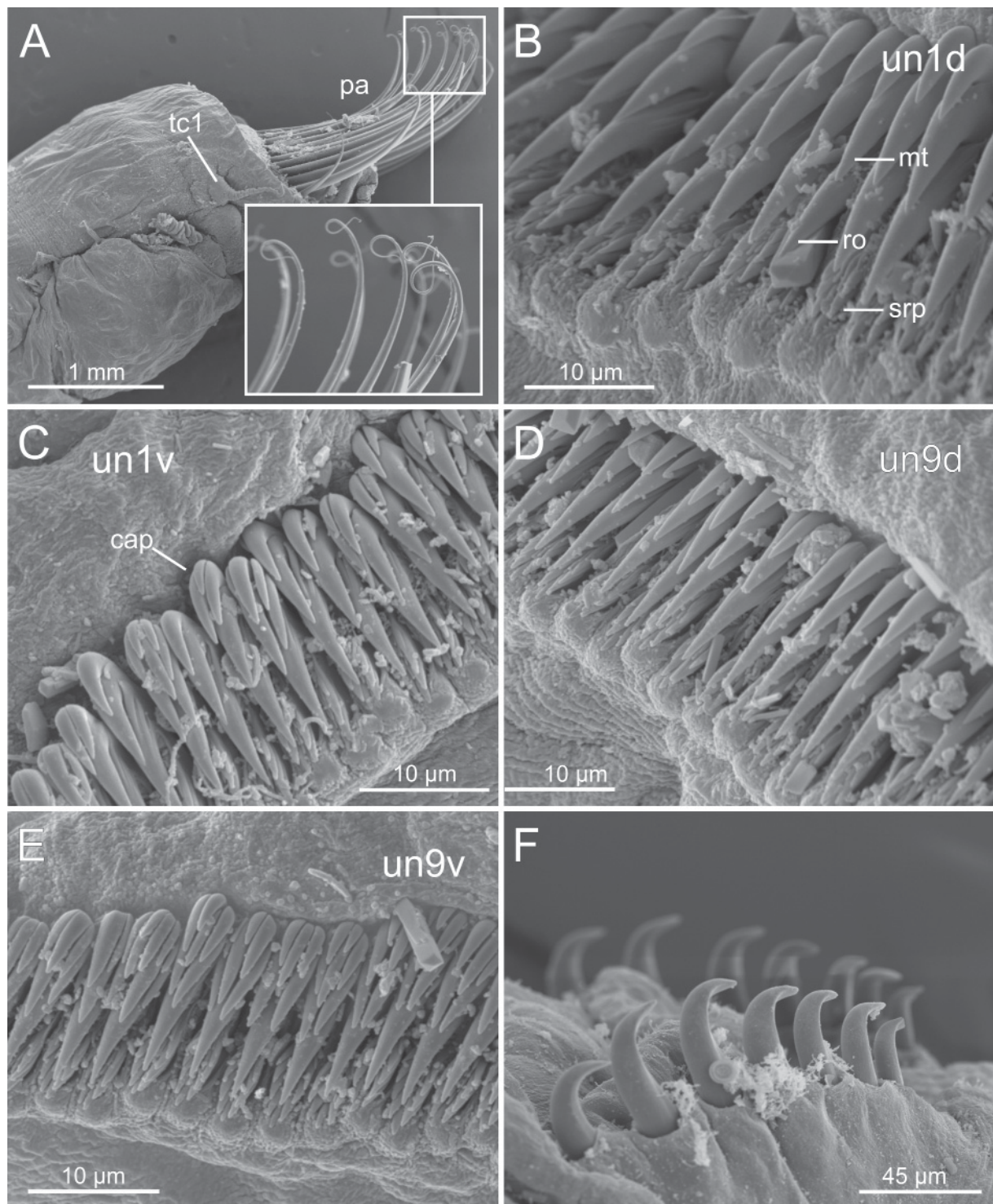


Fig. 12. *Cistenides hyperborea* Malmgren, 1866. SEM micrographs from a medium-sized specimen (BIOICE sample 2660, IINH-40474). **A.** Anterior end, right lateral view (framed: paleae distal end, detail). **B–C.** Unciniger 1, dorsal and ventral uncini, respectively. **D–E.** Unciniger 9, dorsal and ventral uncini, respectively. **F.** Scaphal hooks.

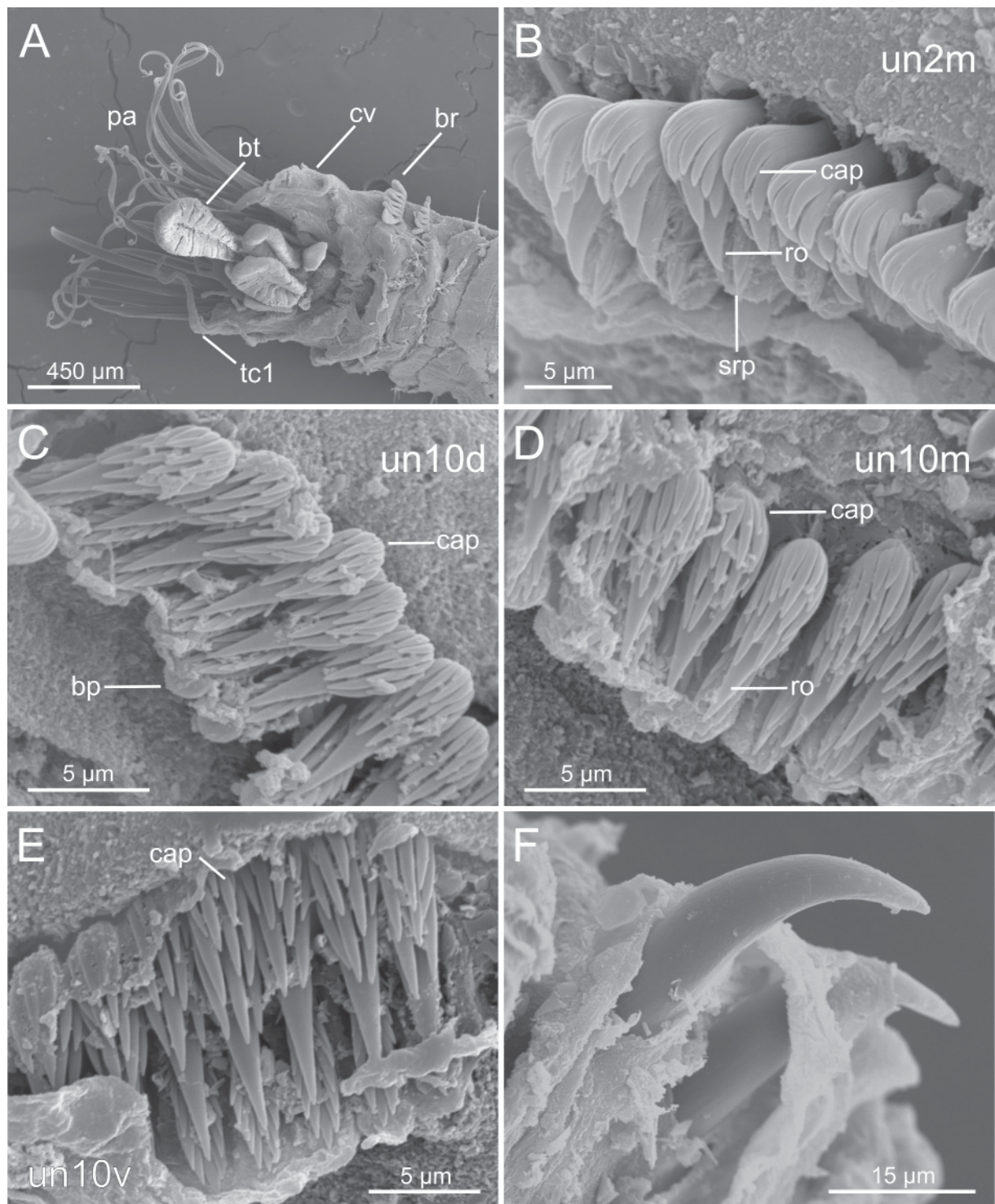


Fig. 13. *Cistenides hyperborea* Malmgren, 1866. SEM micrographs from a small specimen (BIOICE sample 2041, IINH-40470). **A.** Anterior end, ventral view. **B.** Unciniger 2, median uncini. **C–E.** Unciniger 10, dorsal, median and ventral uncini, respectively. **F.** Scaphal hooks.

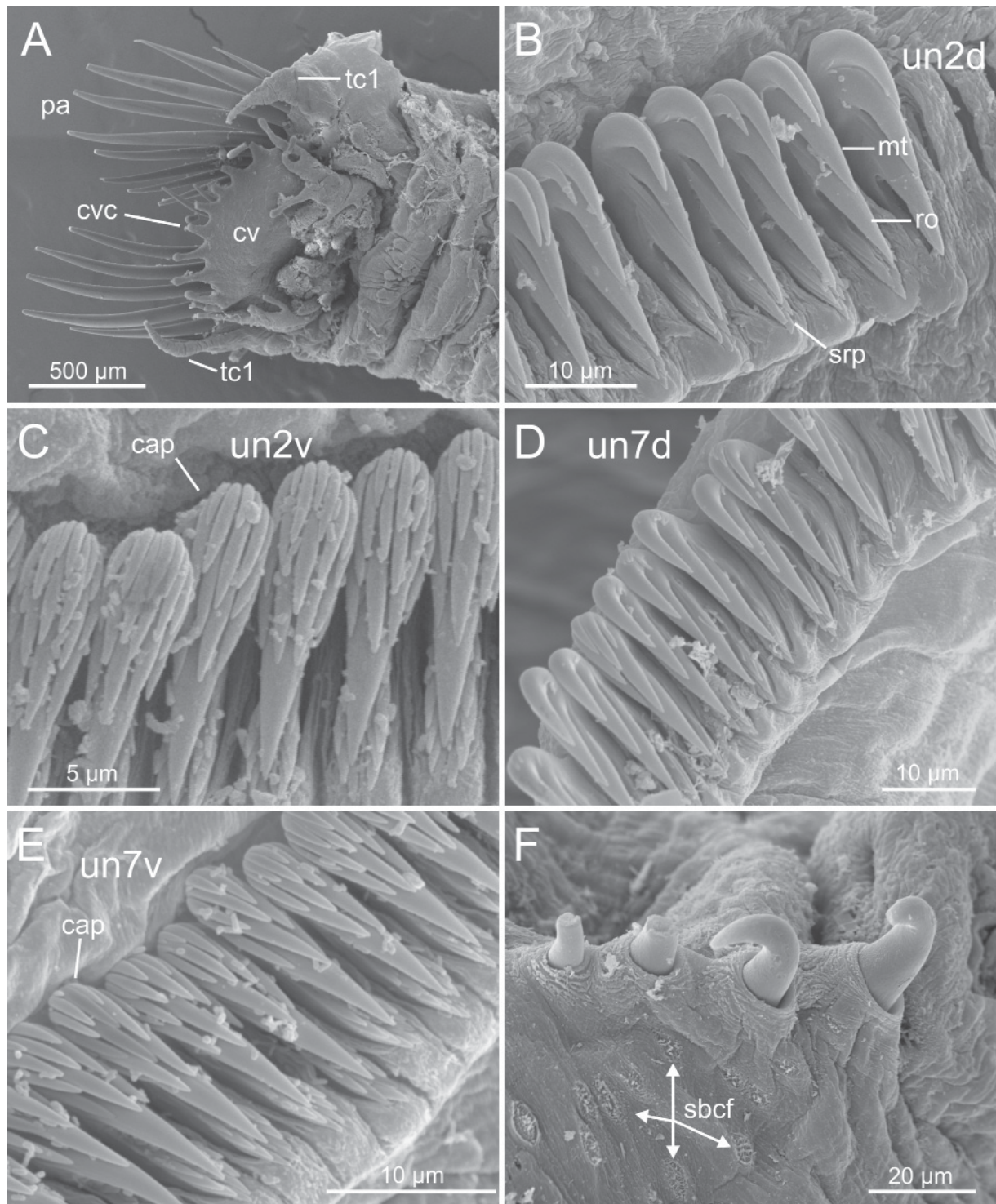


Fig. 14. *Cistenides hyperborea* Malmgren, 1866. SEM micrographs from two medium-sized specimens (BIOICE sample 2060, IINH-40471). **A.** Anterior end, ventral view. **B–C.** Unciniger 2, dorsal and ventral uncini, respectively. **D–E.** Unciniger 7, dorsal and ventral uncini, respectively. **F.** Scaphal hooks and scaphal basis ciliary patches.

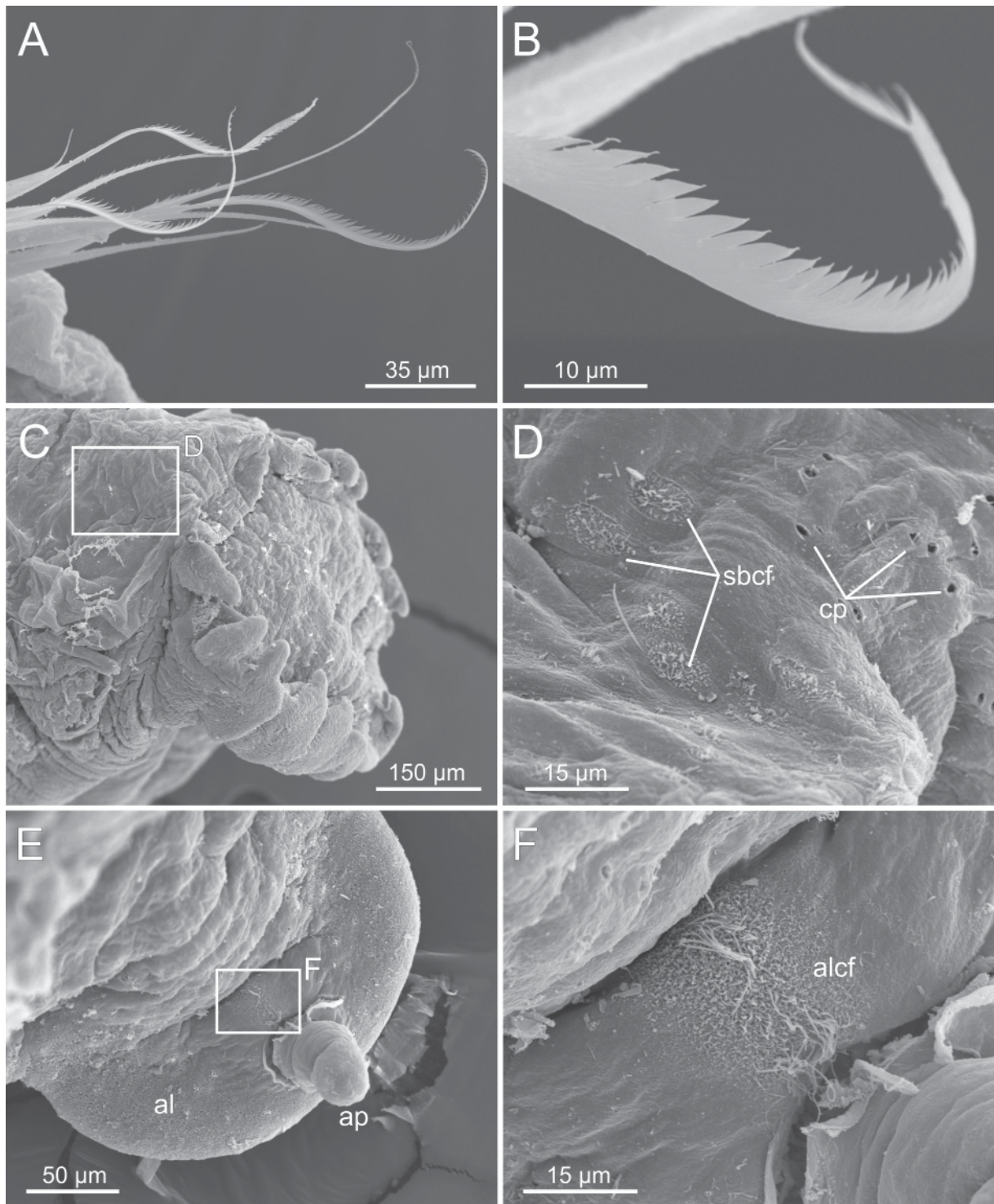


Fig. 15. *Cistenides hyperborea* Malmgren, 1866. SEM micrographs from two medium-sized specimens (BIOICE sample 2060, IINH-40471). **A.** Mid-body parapodium, notochaetae. **B.** Notochaeta, serrated distal end, detail. **C.** Scaphe, dorso-lateral view. **D.** Cuticular structures (framed in C). **E.** Anal lobe and anal papilla. **F.** Ciliary field dorsal to anal lobe (framed in E).

found in the genus (Fig. 12B, D). This pattern of variation is more evident in small specimens (sample 2041, Fig. 13), where the dorsal uncini are also of avicular type (Fig. 13B–D), similar to ventral ones (Fig. 13E). Therefore, these results suggest that the BIOICE specimens of *C. hyperborea* show changes in the uncini shape during ontogenetic development and that the uncini type varies accordingly depending on the age of the specimen. This variation, as explained above, occurs within a torus and along the body; thus, as small specimens with the avicular type of uncini grow, and uncini are being replaced along the torus, there is a progressive decrease in the number of upper teeth and a loss of the rostrum finally resulting in a *pectinate* type with the typical formula of the genus (i.e., MF:1:1:1).

Specimens from sample 2060 (Fig. 2D) found in a shallow bottom (48 m) in North Iceland, were also identified as *C. hyperborea*. Nevertheless, they show thicker paleae (cf. Fig. 14 vs Fig. 13) than similar-sized specimens from other samples, and dorsal uncini are similar to those of larger specimens found elsewhere (Fig. 14B, D), while ventral ones (Fig. 14C, E) correspond to those of specimens of their size such as those of sample 2660 (Fig. 12C, E). Finally, scaphe hooks appear more strongly curved than would be expected for this species (cf. Fig. 14F vs Figs 11E, 12F, 13F). Therefore, this variation suggests that specimens from sample 2060 may correspond to another species (probably undescribed yet) within a potential species complex present in NE Atlantic waters, that should be assessed in future in combination with molecular studies.

Finally, SEM micrographs show that the tip of the notochaetae is serrated (Fig. 15A–B), and reveal the presence of perforations in the cuticle of the dorsal region anterior to the scaphe (Fig. 15C–D) and of two types of ciliated fields: 1) rounded fields located before the scaphe and among the cuticular perforations (Fig. 15C–D) and 2) one large field located dorsal to the anal lobe and anterior to the anal papilla (Fig. 15E–F). These fields may be related to water irrigation, but certainly a future in-depth study is necessary.

Genus *Lagis* Malmgren, 1866

Type species and type locality

Lagis koreni Malmgren, 1866. Finnmark (Norway), lectotype in Naturhistoriska Riksmuseet, Stockholm, designated by Nilsen *et al.* (1977) (see Holthe 1986).

Remarks

The diagnosis of the genus by Hutchings & Peart (2002) states that neurochaetal uncini have large teeth arranged in two or more rows. Examination of BIOICE specimens of *L. koreni* shows again that there are changes in teeth shape with increasing size of an individual, reflected in a progressive reduction in the number of teeth rows with the age of the animal. This still fits the diagnosis of the genus (“two or more rows”) but a better definition of uncini shape is needed. In contrast to the other species studied here, the uncini are, in all body sizes studied, of the *pectinate* type (see below).

Lagis koreni Malmgren, 1866 Figs 1B, 2B, 3B, 4I–L, 16–17

Lagis koreni Malmgren, 1866: 360.

Pectinaria koreni – Hessle 1917: 80.

Pectinaria (Lagis) koreni – Fauvel 1927: 221. — Holthe 1986: 18. — Hartmann-Schröder 1996: 483.
— Kirkegaard 1996: 283.

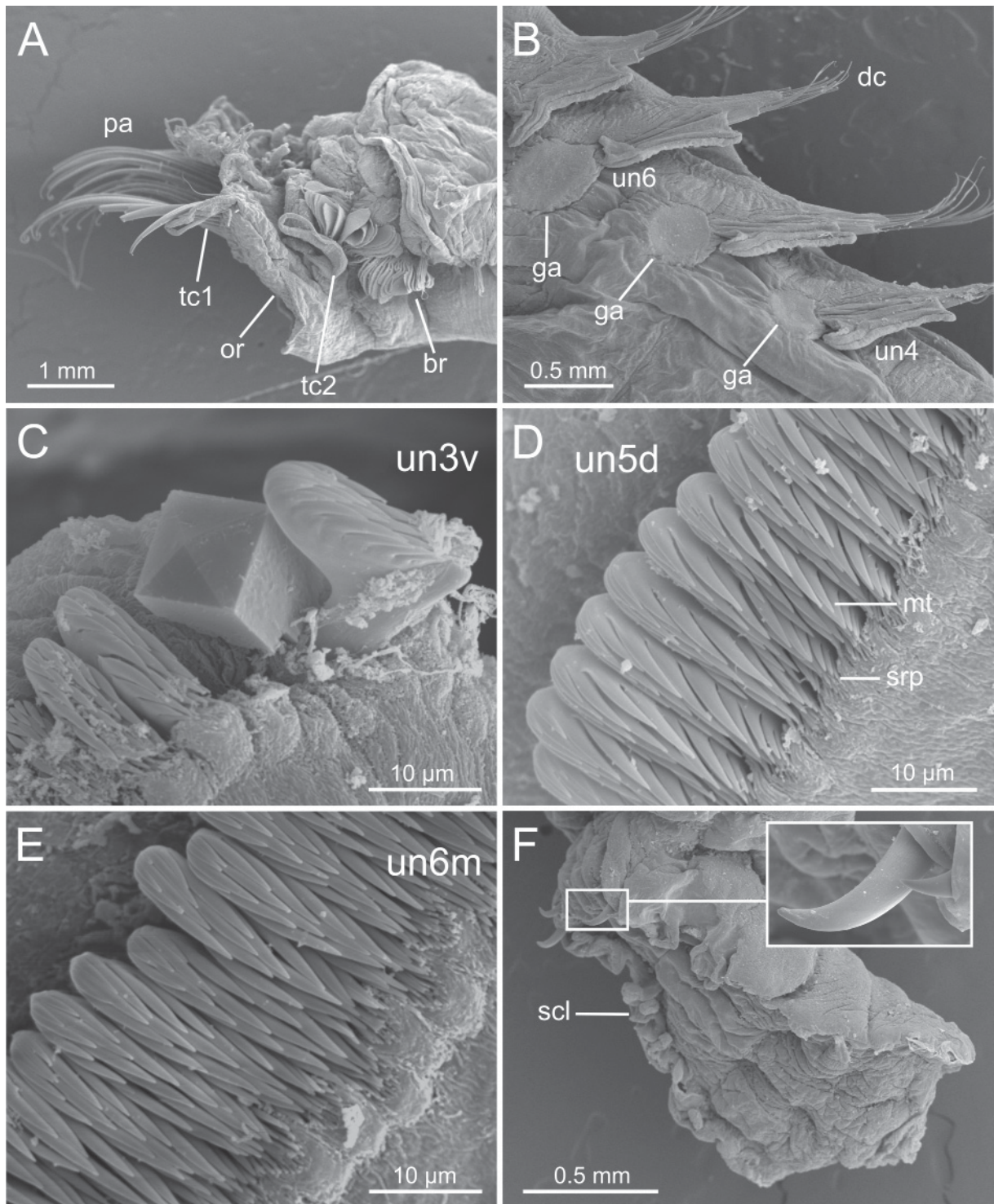


Fig. 16. *Lagis koreni* Malmgren, 1866. SEM micrographs from two specimens: large (A, F) and medium sized (B–E), respectively (BIOICE sample 2827, IINH-40485). **A.** Anterior end, right lateral view. **B.** Uncinigers 4 to 6, ventrolateral view. **C.** Unciniger 3, ventral uncini. **D.** Unciniger 5, dorsal uncini. **E.** Unciniger 6, median uncini. **F.** Scaphe and scaphal hooks (framed: scaphal hook, detail).

Material examined (801 specimens, 75.1% of total specimens identified, in ten samples)

ICELAND – **North Western Fjords** • 1 spec.; BIOICE station 31, sample 2557; 66°21'90" N, 23°04'93" W; 14 Jul. 1993; 7.23°C; 34.34 ppm; 20 m depth; sand; IINH-40483. – **South coast** • 1 spec.; BIOICE station 553, sample 2388; 63°30'10" N, 22°03'70" W; 30 Jun. 1993; 7.13°C; 35.1 ppm; 171 m depth; silty sand; IINH-40478 • 1 spec.; BIOICE station 580, sample 2454; 63°20'10" N, 21°10'20" W; 4 Jul. 1993; 7.19°C; 35.08 ppm; 152 m depth; sediment unknown; MNCN 16.01/18034 • 3 specs; BIOICE station 582, sample 2459; 63°29'80" N, 21°39'40" W; 4 Jul. 1993; 7.10°C; 35.08 ppm; 125 m depth; silt; IINH-40480 • 3 specs; BIOICE station 583, sample 2463; 63°25'40" N, 21°39'89" W; 5 Jul. 1993; 7.12°C; 35.08 ppm; 133 m depth; silty sand and gravel; IINH-40481 • 6 specs; BIOICE station 588, sample 2477; 63°39'69" N, 20°49'63" W; 6 Jul. 1993; 7.80°C; 34.92 ppm; 64 m depth; shelly sand; IINH-40482 • 4 specs; BIOICE station 715, sample 2818; 63°14'64" N, 17°50'70" W; 25 Aug. 1995; 7.18°C; 35.14 ppm; 206 m depth; silty sand; IINH-40484 • 770 specs; BIOICE station 718, sample 2827; 63°41'00" N, 17°39'90" W; 25 Aug. 1995; 8.79°C; 34.77 ppm; 44 m depth; sandy silt; IINH-40485 • 7 specs; BIOICE station 309, sample 3078; 63°28'16" N, 19°33'18" W; 13 Jul. 1997; 9.55°C; 33.97 ppm; 21 m depth; sediment unknown; IINH-40486 • 5 specs; BIOICE station 726, sample 3257; 63°20'10" N, 19°52'20" W; 11 Sep. 2001; 7.92°C; 35.19 ppm; 221 m depth; sediment unknown; IINH-40487.

Occurrence

Off south-western coast of Iceland (Fig. 1B). Depth range: 20 to 221 m; bottom temperature range: 7.10 to 9.55°C (Fig. 2B). The shallowest sample (2557; 20 m) was found in north-western fjords. The easternmost sample (2827; 44 m) contains the highest number of pectinariid specimens found in any BIOICE sample (770; 96.5%). Water mass/es: MNAW.

Remarks

Body and chaetal features match well those of *L. koreni* provided by Holthe (1986), including the shape and number of paleae (Figs 4I, K, 16A, 17A), smooth opercular rim (Fig. 4I, K), cirrate cephalic veil (Fig. 17A–B), a crenulated scaphe (Fig. 16F) provided with slightly distally curved hooks (Figs 16F insert, 17F), and 16 uniramous and 12 biramous segments (see Holthe 1986). However, SEM examination again reveals differences between the uncini features and those reported in the literature, that seem related to morphological variability according to size/age. Large and mid-sized specimens (Fig. 16C–E) bear uncini as described in the literature, i.e., having 3–4 vertical rows with many teeth (6–8) of similar size each and showing no variation across the body or within an individual torus; by contrast, uncini of small specimens (Fig. 17C–E) have more rows of teeth (5–7) in frontal view (Fig. 16E vs Fig. 17D–E).

Finally, SEM micrographs also show circular areas under the parapodia with segmentary arrangement (Fig. 16B); these are likely nephridial or glandular and may have a role in tube building, although this should be confirmed in future morphological studies.

Key to NE Atlantic species of *Pectinariidae*

The key below is based on Gil (2011), who, in turn, adapted those from Fauchald (1977), Holthe (1986) and Hutchings & Peart (2002), and is also complemented with recent papers such as Nishi *et al.* (2014), Zhang & Qiu (2017), Nogueira *et al.* (2019) and Zhang *et al.* (2019).

Several features concerning morphological changes of uncini related to growth as commented above are included. A formula confronting the number of chaetigers with only notopodia or dorsal chaetae (dc) vs those with biramous parapodia or also with ventral uncini (vu) is included as a key character for discriminating between genera. This is referred mostly for Icelandic specimens because of the many inconsistencies found in the literature across species (see Discussion).

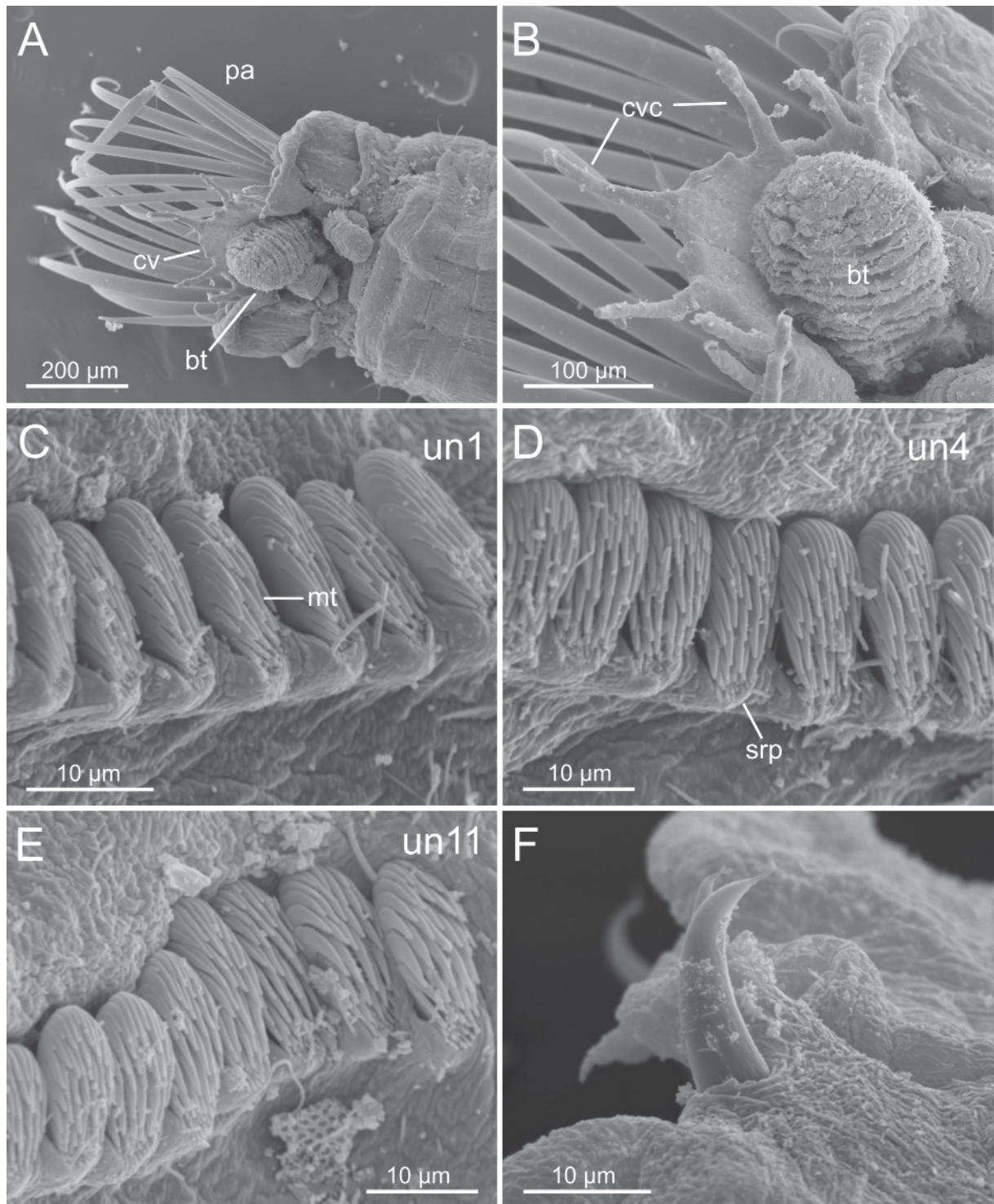


Fig. 17. *Lagis koreni* Malmgren, 1866. SEM micrographs of a small specimen (BIOICE sample 3078, IINH-40486). **A.** Anterior end, ventral view. **B.** Cephalic veil, detail. **C–E.** Uncinigers 1, 4 and 11, uncini. **F.** Scaphal hook.

1. Cephalic veil (tentacular membrane) smooth; scaphe indistinctly separated from abdomen; dc/vu = 17/14 (*Petta*) *Petta pusilla* Malmgren, 1866*
 – Cephalic veil cirrate; scaphe distinctly separated from abdomen; dc/vu not 17/14 2
2. Opercular rim with cirri (serrated) or lappets; dc/vu = 17/13 (*Amphictene*)
 *Amphictene auricoma* (O.F. Müller, 1776)
 – Opercular rim smooth; dc/vu = 17/13 or different 3
3. Cephalic veil attached to lateral margin of opercular rim; dc/vu = 16/12 (*Lagis*)
 *Lagis koreni* Malmgren, 1866
 – Cephalic veil free from opercular rim; dc/vu not 16/12 4
4. Uncini with more than one longitudinal row of major teeth in well-developed specimens; dc/vu = 17/13 (*Pectinaria*) *Pectinaria belgica* (Pallas, 1766)*
 – Uncini with only one longitudinal row of major teeth in well-developed specimens; dc/vu = 17/12 (*Cistenides*) 5
5. Paleae short, with blunt tips; all uncini of *pectinate* type, with one vertical row of teeth, no dorsoventral variation in same unciniger or along the body *Cistenides granulata* (Linnaeus, 1767)
 – Paleae long, with pointed tips; uncini shape *avicular* type in small and medium-sized specimens showing dorso-ventral variation; *pectinate* type (single vertical row of teeth) within the same chaetiger and along the body in large ones *Cistenides hyperborean* Malmgren, 1866

* Species not found in BIOICE samples.

Discussion

Hutchings & Peart (2002), in their revision of the Australian Pectinariidae, stated that “Individuals belonging to the Pectinariidae are easy to recognise, but the systematics of this widely distributed family has been neglected. The confused identity of the common European species suggests that many species may have been misidentified or perhaps several species confused under one name”. This is also true for the NE Atlantic fauna, where the most recent monograph was provided by Holthe (1986); this contrasts with other areas such as China, Brazil and Australia where knowledge on the family has substantially been improved recently (e.g., Sun & Qiu 2012; Zhang & Qiu 2017; Zhang *et al.* 2015, 2019; Nogueira *et al.* 2019).

Abundance, geographical and bathymetric distribution

Among the BIOICE material, *L. koreni* was the most abundant species (> 75%) followed by *A. auricoma* and then by the two species of *Cistenides*. These results agree with those in previous works for the NE Atlantic (e.g., Holthe 1986; Hartmann-Schröder 1996; Gil 2011). Each species mostly shows a homogenous geographic and bathymetric distribution including water temperature, but species of *Cistenides* show slightly wider ranges; the latter are mostly distributed in northern, colder water masses (Figs 1B–C, 2B–C) while *A. auricoma* and *L. koreni* are found in warmer Atlantic waters (see Hansen & Østerhus 2000; Brix & Svavarsson 2010; Meißner *et al.* 2014).

Uncinal morphology

The Terebellomorpha and other tubicolous polychaetes have hook-like neurochaetae (i.e., uncini) that allow for the movement of the worm inside the tube (Woodin & Merz 1987; Merz & Woodin 2000). Previous works such as Hutchings & Glasby (1988), Glasby & Glasby (2006) and Nogueira *et al.* (2010) reported on the variation of their shape during ontogeny in several genera of Terebellidae (e.g., *Polycirrus* Grube, 1850 and *Loimia* Malmgren, 1866). Thus, uncini of fully-grown specimens bear a

vertical series of teeth while in young ones there is a large number arranged in transverse series. The aforementioned authors also affirm that the extent of this variability, both within the same neuropodial row and between segments along the body, has never been widely appreciated, although Duchène & Bhaud (1988) previously reported similar patterns in six terebellid species. Because of that, and its apparent correlation with other relevant taxonomic characters, Glasby & Glasby (2006) proposed to consider these variations in future phylogenetic studies of the group. Garraffoni & Camargo (2006) also stated that Terebellidae uncini "... is a rich source of taxonomic information that can be used to help assess evolutionary relationships within the family". Our study demonstrates that this approach should be extended to the Pectinariidae in general, and especially to the genus *Cistenides*.

Recently, Nogueira *et al.* (2019), following Hutchings & Peart (2002), considered the uncini of *Cistenides* as being "pectinate, with single longitudinal row of teeth" and reported for the first time in the family the presence of avicular uncini when diagnosing the genus *Petta*: "the uncini are breviavicular rather than bipectinate (sensu Holthe 1986), with transverse series of progressively shorter distal wards secondary teeth on top of the main fang." (Nogueira *et al.* 2019: 491). Our observations suggest that: 1) in *Cistenides* the uncini type change as the specimen grows, and this should be considered in the diagnosis of the genus; 2) the presence of avicular uncini in *Petta* should be also confirmed in small specimens; 3) avicular uncini are not exclusive of *Petta* but are also present in Icelandic specimens of *C. hyperborea* as demonstrated here for young specimens of the latter. Nogueira *et al.* (2019) also pointed out that SEM examination is needed to truly assess the uncini features, by comparing SEM micrographs of *Petta alissoni* Nogueira *et al.*, 2019 with those obtained with light microscope (cf. Nogueira *et al.*, 2019: fig. 11d vs fig. 12f–g). Thus, images obtained with the latter show similar uncini to those described and illustrated by Holthe (1986: fig. 7f) as having a "rostrum surmounted by a large tooth", but the use of SEM demonstrated that this is a wrong interpretation of the actual uncini shape. Unfortunately, specimens of *Petta pusilla* Malmgren, 1866, the type species of the genus, were not found among the BIOICE material in order to test this hypothesis and therefore future work is needed to address this.

Dorsal chaetae versus ventral uncini (dc/vu)

Again, following Hutchings & Peart (2002), some generic characters seem to overlap in Pectinariidae, and therefore it seems advisable to find additional diagnostic elements. In fact, these authors propose to include the number of uniramous vs biramous segments, i.e., the number of chaetigers with dorsal chaetae (dc) vs those with ventral uncini (vu), suggesting the following values for each genus: *Amphictene* (17/13), *Cistenides* (17/13), *Lagis* (16/12), *Pectinaria* (17/13) and *Petta* (18/14), although they recognize that the validity of this character needs further testing because some species have reduced posterior notopodia that may easily be overlooked. The study of BIOICE specimens of *Cistenides* confirms the actual number of chaetigers with thoracic notochaetae (17), but we also suggest to modify the number of neuropodial uncinigerous tori from 12 to 13 (see below).

Other taxonomical characters

Hutchings & Peart (2002) include the character "notochaetal ornamentation" among those most relevant for Pectinariidae genera; in the case of *Cistenides*, this feature was not described. Our examination of *C. granulata* and *C. hyperborea* reveals that the distal half of the notochaetae is clearly serrated and this should be added to the genus diagnosis. The same authors also stated that the actual size of the opercular paleae may not be diagnostic because they may be worn by digging and can presumably be replaced during life. From our examination of BIOICE specimens we think that, even considering that the distal end may be worn, other characters remain unchanged during development, including length, width or shape of distal end of paleae. In fact, the paleal general shape is proposed here as a reliable cue to recognize potential different species of *C. hyperborea*.

Delimitation of genera

Following Hutchings & Peart (2002), the characters traditionally used to separate the pectinariid genera are: 1) the degree of fusion of the cephalic veil to the operculum; 2) the marginal ornamentation of the cephalic veil and the operculum; 3) the number of uncinigerous segments; 4) the number of vertical rows of teeth on the uncini, and 5) the degree of separation of the scaphe from the abdomen. As we noted above, Hutchings & Peart (2002) also included details of the notochaetae for the genera represented in Australian waters.

Hutchings & Peart (2002) diagnosed the genus *Cistenides* as having 13 uncinigers and proposed to transfer several species to *Pectinaria* because of having two rows of teeth in uncini instead of one. Following Gil (2011), the genus *Cistenides* as erected by Malmgren (1866) was originally diagnosed as having 12 uncinigers; this diagnosis was followed by many authors such as Fauvel (1927) and Annenkova (1929) and is also followed here. Gil (2011) revised the descriptions of the species attributed by Hutchings & Peart (2002) to this genus (some described as having 13 uncinigers while others with 12), and considered that the genus should be referred as presenting 12–13 uncinigers; however, Gil (2011) also provided a key for pectinariid genera where *Cistenides* is characterized as having 12 uncinigers and *Pectinaria* 13 because this condition was found in specimens of both European species of this genus (both present in BIOICE material), one of them being *C. granulata*, the type species of the genus.

Gil (2011) agrees with Hutchings & Peart (2002) in considering the number of rows of teeth in the uncini as a relevant diagnostic character at the genus level and thus uncini of *Cistenides* would have large teeth arranged in 1–2 rows. In this context, we suggest that this character only corresponds to large specimens and until the ontogeny of all species of Pectinariidae is known, the shape of the uncini should be used with caution when separating genera and species.

It is likely that an in-depth revision of the family in the NE Atlantic, including a molecular approach, may reveal a higher diversity. For instance, Jolly *et al.* (2005) observed different patterns of genetic structure in different populations for *L. koreni* along Brittany and the English Channel; they found a “surprisingly deep phylogeographic break” which was associated with a biogeographic boundary along the western coast of Brittany and suggested the occurrence of potential cryptic or sibling species. In the same vein, Nygren *et al.* (2018) demonstrated the presence of more than 25 species of the genus *Terebellides* Sars, 1835 (Trichobranchidae) along the NE Atlantic coast.

Acknowledgements

The senior author wishes to thank Gudmundur Gudmundsson, Director of Collections and Systematics Department, Náttúrufræðistofnun Íslands (Icelandic Institute of Natural History) for the loan of the specimens. Special thanks to all the people involved in sorting of specimens from the BIOICE samples in the SMC. Ada Castro and Catalina Sueiro (SAIN, UDC) assisted in the preparation of specimens. This study was partly supported by the *Fauna Ibérica* research project (*Polychaeta VII, Palpata, Canalipalpata II* - PGC2018-095851-B-C64) to J. Parapar and J. Moreira. The Bibliographic Service of the Universidade da Coruña is also thanked for providing useful bibliography. Authors would like to thank the reviewers Pat Hutchings and Joao Miguel de Matos Nogueira as well as the topical and desk editors of *EJT* Rudy Jocqué and Kristiaan Hoedemakers for their constructive comments on the manuscript.

References

Annenkova N.P. 1929. Beiträge zur Kenntnis der Polychaeten-Fauna der USSR. I. Fam. Pectinariidae Quatrefages (Amphictenidae Malmgren) und Ampharetidae Malmgren. *Annuaire du Musée zoologique*

de l'Académie des Sciences de l'URSS [Ezhegodnik Zoologicheskogo Museya] AN SSSR 30 (3): 477–502.

Brix S. & Svavarsson J. 2010. Distribution and diversity of desmosomatid and nannoniscid isopods (Crustacea) on the Greenland–Iceland–Faeroe Ridge. *Polar Biology* 33: 515–530.

<https://doi.org/10.1007/s00300-009-0729-8>

Chambers S.J. & Woodham A. 2003. A new species of *Chaetozone* (Polychaeta: Cirratulidae) from the deep water in the northeast Atlantic, with comments on the diversity of the genus in cold northern waters. *Hydrobiologia* 496: 41–48. <https://doi.org/10.1023/A:1026116008735>

Day J.H. 1967. *A Monograph on the Polychaeta of Southern Africa*. British Museum of Natural History Publication 656. Trustees of the British Museum (Natural History), London.

<https://doi.org/10.5962/bhl.title.8596>

Duchène J.C. & Bhaud M. 1988. Uncinial patterns and age determination in terebellid polychaetes. *Marine Ecology Progress Series* 49: 267–275. <https://doi.org/10.3354/meps049267>

Fauchald K. 1977. The polychaete worms. Definitions and keys to the orders, families and genera. *Natural History Museum of Los Angeles County, Science Series* 28: 1–188.

Fauvel P. 1927. Polychètes sédentaires. Addenda aux errantes, archiannélides, myzostomaires. *Faune de France* 16: 1–494.

Garraffoni A.R.S. & Camargo M.G. 2006. First application of morphometrics in a study of variations in uncinial shape present within the Terebellidae (Polychaeta). *Zoological Studies* 45 (1): 75–80.

Gil J. 2011. *The European Fauna of Annelida Polychaeta*. PhD Thesis, Universidade de Lisboa, Portugal.

Glasby C.J. & Glasby T.M. 2006. Two types of uncini in *Polycirrus* (Polychaeta: Terebellidae: Polycirrinae) revealed using geometric morphometrics. *Journal of Natural History* 40 (5–6): 237–253.

<https://doi.org/10.1080/00222930600627137>

Hansen B. & Østerhus S. 2000. North Atlantic–Nordic Seas exchanges. *Progress in Oceanography* 45: 109–208. [https://doi.org/10.1016/S0079-6611\(99\)00052-X](https://doi.org/10.1016/S0079-6611(99)00052-X)

Hartman O. 1941. Polychaetous annelids. Part IV. Pectinariidae. *Allan Hancock Pacific Expeditions* 7: 325–345.

Hartmann-Schröder G. 1971. *Annelida, Borstenwürmer, Polychaeta*. Die Tierwelt Deutschlands 58, 1st ed. Gustav Fischer, Jena.

Hartmann-Schröder G. 1996. *Annelida, Borstenwürmer, Polychaeta*. Die Tierwelt Deutschlands 58, 2nd ed. Gustav Fischer, Jena.

Hessle C. 1917. Zur Kenntnis der terebellomorphen Polychaeten. *Zoologiska Bidrag från Uppsala* 5: 39–258.

Holthe T. 1986. *Polychaeta Terebellomorpha*. Marine Invertebrates of Scandinavia 7, Norwegian University Press, Oslo.

Hutchings P. 2000. Family Pectinariidae. In: Beesley P.L., Ross G.J.B. & Glasby C.J. (eds) *Polychaetes & Allies: The Southern Synthesis. Fauna of Australia. Vol. 4A. Polychaeta, Myzostomida, Pogonophora, Echiura, Sipuncula*: 294–299. CSIRO Publishing, Melbourne.

Hutchings P.A. & Glasby C.J. 1988. The Amphitritinae (Polychaeta: Terebellidae) from Australia. *Records of the Australian Museum* 40 (1): 1–60. <https://doi.org/10.3853/j.0067-1975.40.1988.150>

- Hutchings P.A. & Peart R. 2002. A review of the genera of Pectinariidae (Polychaeta) together with a description of the Australian fauna. *Records of the Australian Museum* 54: 99–127.
<https://doi.org/10.3853/j.0067-1975.54.2002.1356>
- Jirkov I.A. & Leontovich M.K. 2013. Identification keys for Terebellomorpha (Polychaeta) for the Eastern Atlantic and the North Polar basin. *Invertebrate Zoology* 10 (2): 217–243.
<https://doi.org/10.15298/invertzool.10.2.02>
- Jolly M.T., Jollivet D., Gentil F., Thiébaud E. & Viard F. 2005. Sharp genetic break between Atlantic and English Channel populations of the polychaete *Pectinaria koreni*, along the North coast of France. *Heredity* 94: 23–32. <https://doi.org/10.1038/sj.hdy.6800543>
- Kirkegaard J.B. 1996. Havbørsteorme II. Sedentaria. *Danmarks Fauna* 86: 1–451.
- Kirkegaard J.B. 2001. Polychaetes of the families Glyceridae, Goniadidae and Nereididae from the North Atlantic around Iceland. *Sarsia* 86: 13–20. <https://doi.org/10.1080/00364827.2001.10420457>
- Linnaeus C. 1767. *Systema Naturae*. 12th ed. Laurentius Salvius, Stockholm.
- Long C.D. 1973. Pectinariidae (Polychaeta) from Caribbean and associated waters. *Bulletin of Marine Science* 23: 857–874.
- Malmgren A.J. 1865. Nordiska Hafs-Annulater. *Öfversigt af Konglia Vetenskaps-Akademiens Förhandlingar, Stockholm* 21: 51–110, 181–192.
- Malmgren A.J. 1866. Nordiska Hafs-Annulater. *Öfversigt af Konglia Vetenskaps-Akademiens Förhandlingar, Stockholm* 22: 355–410.
- Meißner K., Fiorentino D., Schnurr S., Martínez Arbizu P., Huettmann F., Holst S., Brix S. & Svavarsson J. 2014. Distribution of benthic marine invertebrates at northern latitudes – An evaluation applying multi-algorithm species distribution models. *Journal of Sea Research* 85: 241–254.
<https://doi.org/10.1016/j.seares.2013.05.007>
- Merz R.A. & Woodin S.A. 2000. Hooked setae: tests of the anchor hypothesis. *Invertebrate Biology* 119: 67–82. <https://doi.org/10.1111/j.1744-7410.2000.tb00175.x>
- Moreira J. & Parapar J. 2012. Two new species of *Sphaerodoropsis* Hartman & Fauchald, 1971 (Polychaeta: Sphaerodoridae) from Iceland (BIOICE programme). *Marine Biology Research* 8 (7): 584–593. <https://doi.org/10.1080/17451000.2011.638929>
- Moreira J. & Parapar J. 2015. A new species of *Sphaerodoridium* Lützen, 1961 from Iceland (Polychaeta: Sphaerodoridae). *Zootaxa* 3911 (1): 91–105. <https://doi.org/10.11646/zootaxa.3911.1.5>
- Müller O.F. 1776. *Zoologicae Danicae Prodrum, seu Animalium Daniae et Norvegiae indigenarum characteres, nomina et synonyma imprimis popularium*. Hallageriis, Havniae, Copenhagen.
<https://doi.org/10.5962/bhl.title.13268>
- Nilsen R., Kirkegaard J.B. & Lemche H. 1977. *Pectinaria* Lamarck, 1818 (Polychaeta), and the species names *P. belgica* (Pallas, 1766) and *P. koreni* (Malmgren, 1866) to be validated under the plenary powers. *Bulletin of Zoological Nomenclature* 34: 112–122. <https://doi.org/10.5962/bhl.part.14601>
- Nilsson D. 1928. Neue und alte Amphicteniden. *Göteborgs Kungelige Vetenskaps- och Vitterhets Samhälles Handlingar, Series 4* 33: 1–96.
- Nishi E., Matsuo K., Kazama-Wakabayashi M., Mori A., Tomioka S., Kajihara S., Hamaguchi M., Kajihara M. & Hutchings P. 2014. Partial revision of Japanese Pectinariidae (Annelida: Polychaeta), including redescriptions of poorly known species. *Zootaxa* 3895 (3): 433–445.
<https://doi.org/10.11646/zootaxa.3895.3.8>

- Nogueira J.M.M., Hutchings P.A. & Fukuda M.V. 2010. Morphology of terebelliform polychaetes (Annelida: Polychaeta: Terebelliformia), with a focus on Terebellidae. *Zootaxa* 2460: 1–185. <https://doi.org/10.11646/zootaxa.2460.1.1>
- Nogueira J.M.M., Ribeiro W.M.G., Carrerette O. & Hutchings P. 2019. Pectinariidae (Annelida, Terebelliformia) from off southeastern Brazil, southwestern Atlantic. *Zootaxa* 4571 (4): 489–509. <https://doi.org/10.11646/zootaxa.4571.4.3>
- Nygren A., Parapar J., Pons J., Meißner K., Bakken T., Kongsrud J.A., Oug E., Gaeva D., Sikorski A., Johansen R.A., Hutchings P.A., Lavesque N. & Capa M. 2018. A mega-cryptic species complex hidden among one of the most common annelids in the North East Atlantic. *PLoS One* 13 (6): e0198356. <https://doi.org/10.1371/journal.pone.0198356>
- Parapar J. 2003. Oweniidae (Annelida, Polychaeta) from Icelandic waters, collected by the BIOICE project, with a description of *Myrioglobula islandica* n. sp. *Sarsia* 88: 274–290.
- Parapar J. 2006. The genera *Myriochele* and *Myrioglobula* (Polychaeta, Oweniidae) in Icelandic waters with the revision of type material of *Myriochele heeri* Malmgren, 1867, and the description of a new species. *Journal of Natural History* 40 (9–10): 523–547. <https://doi.org/10.1080/00222930600711758>
- Parapar J., Moreira J. & Helgason G.V. 2011a. Distribution and diversity of the Opheliidae (Annelida, Polychaeta) on the continental shelf and slope of Iceland, with a review of the genus *Ophelina* in northeast Atlantic waters and description of two new species. *Organisms, Diversity and Evolution* 11: 83–105. <https://doi.org/10.1007/s13127-011-0046-2>
- Parapar J., Moreira J. & Helgason G.V. 2011b. Taxonomy and distribution of *Terebellides* (Polychaeta, Trichobranchidae) in Icelandic waters, with the description of a new species. *Zootaxa* 2983: 1–20. <https://doi.org/10.11646/zootaxa.2983.1.1>
- Parapar J., Helgason G.V., Jirkov I. & Moreira J. 2011c. Taxonomy and distribution of the genus *Amphicteis* (Polychaeta: Ampharetidae) collected by the BIOICE project in Icelandic waters. *Journal of Natural History* 45 (23–24): 1477–1499. <https://doi.org/10.1080/00222933.2011.558640>
- Parapar J., Moreira J. & Helgason G.V. 2013a. First record of genus *Orbiniella* Day, 1954 (Polychaeta: Orbiniidae) in North Atlantic Ocean with the description of a new species. *Zootaxa* 4006: 330–346. <https://doi.org/10.11646/zootaxa.4006.2.5>
- Parapar J., Helgason G.V., Jirkov I. & Moreira J. 2013b. Polychaetes of the genus *Ampharete* (Polychaeta: Ampharetidae) collected in Icelandic waters during the BIOICE project. *Helgoland Marine Research* 66: 331–344. <https://doi.org/10.1007/s10152-011-0274-z>
- Parapar J., Aguirrezabalaga F. & Moreira J. 2014a. First record of Longosomatidae (Annelida: Polychaeta) from Iceland with a worldwide review of diagnostic characters of the family. *Journal of Natural History* 48 (17–18): 983–998. <https://doi.org/10.1080/00222933.2013.859316>
- Parapar J., Helgason G.V., Jirkov I. & Moreira J. 2014b. Diversity and taxonomy of Ampharetidae (Polychaeta) from Icelandic waters. *Polish Polar Research* 35 (2): 311–340.
- Petersen M.E. 2000. A new genus of Fauveliopsidae (Annelida: Polychaeta), with a review of its species and redescription of some described taxa. *Bulletin of Marine Science* 67: 491–515.
- Pettibone M.H. 1954. Marine Polychaete worms from Point Barrow, Alaska, with additional records from the North Atlantic and North Pacific. *Proceedings of the United States National Museum* 103 (3324): 203–356. <https://doi.org/10.5479/si.00963801.103-3324.203>
- Read G. & Fauchald K. (eds) 2019. World Polychaeta database. Pectinariidae Quatrefages, 1866. Available from <http://www.marinespecies.org/aphia.php?p=taxdetails&id=980> [accessed 17 Mar. 2019].

- Rouse G.V. & Pleijel F. 2001. *Polychaetes*. Oxford University Press, New York.
- Sanfilippo R. 2001. *Bathyvermilia islandica* (Polychaeta, Serpulidae): a new deep-water species from south of Iceland. *Sarsia* 86: 177–182. <https://doi.org/10.1080/00364827.2001.10420473>
- Sigvaldadóttir E. 2002. Polychaetes of the genera *Prionospio* and *Aurospio* (Spionidae, Polychaeta) from Icelandic waters. *Sarsia* 87: 201–215. <https://doi.org/10.1080/00364820260294842>
- Silva T., Gislason A., Astthorsson O.S. & Marteinsdóttir G. 2017. Distribution, maturity and population structure of *Meganyctiphanes norvegica* and *Thysanoessa inermis* around Iceland in spring. *PLoS One* 12 (11): e0187360. <https://doi.org/10.1371/journal.pone.0187360>
- Sun Y. & Qiu J.-W. 2012. A new species of *Lagis* (Polychaeta: Pectinariidae) from Hong Kong. *Zootaxa* 3264: 61–68. <https://doi.org/10.11646/zootaxa.3264.1.4>
- Wesenberg-Lund E. 1951. Polychaeta. *The Zoology of Iceland* 2 (19): 1–181.
- Wolf P.S. 1984. Family Pectinariidae Quatrefages, 1865. Chapter 50. In: Uebelacker J.M. & Johnson P.G. (eds) *Taxonomic Guide to the Polychaetes of the Northern Gulf of Mexico, Volume VII*: 50-1–50-10. U.S. Department of the Interior Minerals Management Service.
- Woodin S.A. & Merz R.A. 1987. Holding on by their hooks: anchors for worms. *Evolution* 41: 427–432. <https://doi.org/10.1111/j.1558-5646.1987.tb05808.x>
- Zamora J.L., Parapar J., Helgason G.V. & Moreira J. 2020. Taxonomy and distribution of Icelandic Fauveliopsidae (Annelida) collected during the BIOICE project. *Journal of Natural History* 53 (47–48): 2951–2954. <https://doi.org/10.1080/00222933.2020.1757170>
- Zhang J. & Qiu J.W. 2017. A new species of *Pectinaria* (Annelida, Pectinariidae), with a key to pectinariids from the South China Sea. *ZooKeys* 683: 139–150. <https://doi.org/10.3897/zookeys.683.12272>
- Zhang J., Zhang Y. & Qiu J.W. 2015. A new species of *Amphictene* (Annelida, Pectinariidae) from the northern South China Sea. *ZooKeys* 545: 27–36. <https://doi.org/10.3897/zookeys.545.6454>
- Zhang J., Hutchings P. & Kupriyanova E. 2019. A revision of the genus *Petta* Malmgren, 1866 (Annelida: Pectinariidae), with two new species from deep waters of southeastern Australia, and comments on phylogeny of the family. *Zootaxa* 4614 (2): 303–330. <https://doi.org/10.11646/zootaxa.4614.2.3>

Manuscript received: 28 November 2019

Manuscript accepted: 17 March 2020

Published on: 15 June 2020

Topic editor: Rudy Jocqué

Desk editor: Kristiaan Hoedemakers

Printed versions of all papers are also deposited in the libraries of the institutes that are members of the *EJT* consortium: Muséum national d'histoire naturelle, Paris, France; Meise Botanic Garden, Belgium; Royal Museum for Central Africa, Tervuren, Belgium; Royal Belgian Institute of Natural Sciences, Brussels, Belgium; Natural History Museum of Denmark, Copenhagen, Denmark; Naturalis Biodiversity Center, Leiden, the Netherlands; Museo Nacional de Ciencias Naturales-CSIC, Madrid, Spain; Real Jardín Botánico de Madrid CSIC, Spain; Zoological Research Museum Alexander Koenig, Bonn, Germany; National Museum, Prague, Czech Republic.

ZOBODAT - www.zobodat.at

Zoologisch-Botanische Datenbank/Zoological-Botanical Database

Digitale Literatur/Digital Literature

Zeitschrift/Journal: [European Journal of Taxonomy](#)

Jahr/Year: 2020

Band/Volume: [0666](#)

Autor(en)/Author(s): Parapar Julio, Palomanes Veronica, Helgason Gudmundur V.,
Moreira Juan

Artikel/Article: [Taxonomy and distribution of Pectinariidae \(Annelida\) from Iceland with a comparative analysis of uncinal morphology 1-32](#)