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Integrative description of *Macrobiotus canaricus* sp. nov. with notes on *M. recens* (Eutardigrada: Macrobiotidae)

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Abstract. In this paper we describe *Macrobiotus canaricus* sp. nov., a new tardigrade species of the *Macrobiotus hufelandi* group from the Canary Islands. Moreover, with the use of DNA sequencing, we confirm that *Macrobiotus recens* Cuénot, 1932 represents the *hufelandi* group, even though eggs laid by this species do not exhibit the typical *hufelandi* group morphology. Our study is based on both classical taxonomic methods that include morphological and morphometric analyses conducted with the use of light and scanning electron microscopy, and on the analysis of nucleotide sequences of four molecular markers (three nuclear: 18S rRNA, 28S rRNA, ITS-2, and one mitochondrial: COI). Our analyses revealed that *M. canaricus* sp. nov. is most similar to *Macrobiotus almadai* Fontoura *et al.*, 2008 from the Archipelago of the Azores, from which it differs by the absence of granulation patches on the external and internal surfaces of legs I–III as well as by the absence of a cuticular pore in the centre of the external patch on legs I–III. Molecular sequences allowed us to pinpoint the phylogenetic positions of *M. canaricus* sp. nov. and *M. recens* within the *M. hufelandi* group.

Keywords. Canary Islands, COI, *Macrobiotus hufelandi* group, phylogeny, tardigrades.

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Introduction

Tardigrades are a phylum of globally distributed microinvertebrates which inhabit terrestrial and marine habitats (Nelson *et al.* 2015). However, the majority of the over 1200 known species were found and described from terrestrial mosses and lichens (Guidetti & Bertolani 2005; Degma & Guidetti 2007; Degma *et al.* 2009–2017). Although tardigrade faunistic studies have been conducted for more than two centuries, many regions are still very poorly known in terms of tardigrade species composition. One such place is the Canary Islands (Spain), where only two studies on tardigrades

have been performed until now (Heinis 1908; Guil & Guidetti 2005). In total, 13 taxa identified to species level have been reported from the Canary Islands: *Echiniscus arctomys* Ehrenberg, 1853, *E. canadensis* Murray, 1910, *E. quadrispinosus* Richters, 1902, *E. mediantus* Marcus, 1930, *E. trisetosus* Cuénot, 1932, *Isohypsibius tuberculatus* (Plate, 1888), *Macrobiotus echinogenitus* Richters, 1904, *M. furcatus* Ehrenberg, 1859, *M. hufelandi* C.A.S. Schultze, 1834, *M. occidentalis* Murray, 1910, *Milnesium tardigradum* Doyère, 1840, *Minibiotus intermedius* (Plate, 1888) and *M. gumersindoi* Guil & Guidetti, 2005. Moreover, both studies reported undetermined species of the *Macrobiotus hufelandi* group and Heinis (1908) noted an undetermined species of *Echiniscus* C.A.S. Schultze, 1840. However, not all of the thirteen species identifications can be considered certain according to modern taxonomic standards. For example, records of *M. hufelandi* and *M. tardigradum* predate the redescriptions of these species (Bertolani & Rebecchi 1993 and Michalczyk *et al.* 2012, respectively), and thus they need to be verified against modern literature.

We examined moss and lichen samples from the island of Gran Canaria. In two of them, we found two species of the *Macrobiotus hufelandi* group, one new to science and the other conforming to the description of *Macrobiotus recens* Cuénot, 1932. For both these populations, we provide taxonomic descriptions based on morphological and morphometric data acquired with phase contrast light microscopy (PCM) as well as from scanning electron microscopy (SEM). Additionally, we sequenced four DNA markers, three nuclear (18S rRNA, 28S rRNA, ITS-2) and one mitochondrial (COI), that allowed us to pinpoint the phylogenetic positions of the two species. Specimens of *M. cf. recens* conform to the diagnosis of the *hufelandi* group, but they lay eggs that depart from the typical *hufelandi* group egg ornamentation, i.e., instead of inverted goblet-shaped processes, the eggs are equipped with conical processes. Thus, the species either has not (Biserov 1990a, 1990b) or has (Bertolani & Rebecchi 1993; Kaczmarek & Michalczyk 2017b) been classified within the *hufelandi* group. Therefore, the first DNA sequences for *M. recens* allowed us to verify whether the species represents the *hufelandi* group.

Material and methods

Sample processing and tardigrade culturing

The moss and lichen samples containing the new species and *M. cf. recens*, respectively, were collected from two localities in Gran Canaria by Marta Kapała on 16 February 2017 (see Table 1 for details). Samples were examined for terrestrial tardigrades using standard methods described by Dastych (1980) with modification by Stec *et al.* (2015). A total of 21 individuals of the new species and 29 individuals of *M. cf. recens* were extracted from the first and the second sample, respectively. Live animals were placed in a culture to obtain more individuals and eggs required for integrative analyses (see Table 1 for details). Animals were reared on plastic Petri dishes according to the protocol in Stec *et al.* (2015).

Microscopy and imaging

Specimens for light microscopy were mounted on microscope slides in a small drop of Hoyer's medium and secured with a cover slip, following the protocol in Morek *et al.* (2016b). Slides were then dried for five to seven days at 60 °C. Dried slides were sealed with a transparent nail polish and examined under a Nikon Eclipse 50i phase contrast light microscope (PCM) associated with a Nikon Digital Sight DS-L2 digital camera. In order to obtain clean and extended specimens for SEM, tardigrades were processed according to the protocol in Stec *et al.* (2015). In short, specimens were first subjected to a 60 °C water bath for 30 min to obtain fully extended animals, followed by a water/ethanol and an ethanol/acetone series, and then CO₂ critical point drying and finally sputter coated with a thin layer of gold. Specimens were examined under high vacuum in a Versa 3D DualBeam Scanning Electron Microscope at the ATOMIN facility of the Jagiellonian University, Kraków, Poland. Both studied populations were also examined for the presence of males with aceto-orcein staining (Bertolani 1971) in accordance with Stec *et al.* (2016a).

Table 1. Sample collection details and numbers of individuals and eggs used for integrative analyses of *M. canaricus* sp. nov. and *M. cf. recens* Cuénot, 1932. (PCM = phase contrast microscopy; SEM = scanning electron microscopy; DNA = DNA sequencing; ORC = orcein staining)

*Eggs were taken from the culture and incubated individually until hatching, then the juveniles were used for DNA sequencing whereas egg chorions (isogenophores) were mounted on microscope slides in Hoyer's medium.

Species	Sample code	Coordinates; altitude	Sample type, habitat	PCM	SEM	DNA	ORC
<i>M. canaricus</i> sp. nov.	ES.004	28°03'05" N, 15°38'21" W; 1050 m a.s.l.	moss on a tree trunk in a pine forest	120 ind. 42 eggs	20 ind. 15 eggs	4 ind. 3 eggs*	15 ind. 0 eggs
<i>M. cf. recens</i>	ES.006	28°02'38" N, 15°40'22" W; 885 m a.s.l.	lichen on a stone wall	87 ind. 38 eggs	8 ind. 10 eggs	0 ind. 4 eggs*	15 ind. 0 eggs
<i>M. macrocalix</i>	PL.110	49°31'27" N, 21°02'23" E; 254 m a.s.l.	moss on a roof	60 ind. 72 eggs	0 ind. 0 eggs	4 ind. 0 eggs	0 ind. 0 eggs

All figures were assembled in Corel Photo-Paint X6, v. 16.4.1.1281. For structures that could not be satisfactorily focused in a single photograph, a stack of 2–8 images was taken with an equidistance of ca 0.2 µm and assembled manually into a single deep-focus image.

Morphometrics and morphological nomenclature

All measurements are given in micrometres (µm). Sample size was adjusted following recommendations by Stec *et al.* (2016b). Structures were measured only if their orientation was suitable. Body length was measured from the anterior extremity to the end of the body, excluding the hind legs. The terminology used to describe oral cavity armature and egg shell morphology follows Michalczyk & Kaczmarek (2003) and Kaczmarek & Michalczyk (2017b). Macroplacoid length sequence is given according to Kaczmarek *et al.* (2014). Buccal tube length and the level of the stylet support insertion point were measured according to Pilato (1981). The *pt* index is the ratio of the length of a given structure to the length of the buccal tube expressed as a percentage (Pilato 1981). All other measurements and nomenclature are given in accordance with Kaczmarek & Michalczyk (2017b). In brief, the buccal tube width was measured as the external and internal diameter at the level of the stylet support insertion point. Lengths of the claw branches were measured from the base of the claw (i.e., excluding the lunula) to the top of the branch, including accessory points. Distance between egg processes was measured as the shortest line connecting base edges of the two closest processes. Morphometric data were handled using the 'Parachela' v. 1.3 template available from the Tardigrada Register (Michalczyk & Kaczmarek 2013). Tardigrade taxonomy follows Bertolani *et al.* (2014).

Comparative material

The taxonomic key for the *hufelandi* group provided by Kaczmarek & Michalczyk (2017b) was used to determine whether the isolated species had previously been described. After one species could not be identified with the key, we compared it with the original description of the most similar species of the *hufelandi* group, *Macrobiotus almadai* Fontoura *et al.*, 2008. Additionally, we also compared our new species with three paratypes and one egg of *M. almadai* kindly loaned to us by Professor Paulo Fontoura. Moreover, thanks to the courtesy of Matteo Vecchi, we analysed photomicrographs of animals and eggs identified as '*M. recens*' by Maucci (1979) from several Portuguese populations deposited in the Maucci collection. Although our population conformed to the original description of *M. recens*, the description

Table 2. Primers and references for PCR protocols for amplification of the four DNA fragments sequenced in the study.

DNA fragment	Primer name	Primer direction	Primer sequence (5'-3')	Primer source	PCR programme
18S rRNA	18S_Tar_Ff1	forward	AGGCGAAACCGCGAATGGCTC	Stec <i>et al.</i> (2017a)	Zeller (2010)
	18S_Tar_Rr1	reverse	GCCGCAGGCTCCACTCCTGG		
28S rRNA	28SF0001	forward	ACCCVCYNAATTTAAGCATAT	Mironov <i>et al.</i> (2012)	Mironov <i>et al.</i> (2012)
	28SR0990	reverse	CCTTGGTCCGTGTTTCAAGAC		
ITS-2	ITS2_Eutar_Ff	forward	CGTAACGTGAATTGCAGGAC	Stec <i>et al.</i> (2018c)	Stec <i>et al.</i> (2018c)
	ITS2_Eutar_Rr	reverse	TCCTCCGCTTATTGATATGC		
COI	LCO1490	forward	GGTCAACAAATCATAAAGATATTGG	Folmer <i>et al.</i> (1994)	Michalczyk <i>et al.</i> (2012)
	HCO2198	reverse	TAAACTTCAGGGTGACCAAAAAATCA		

is outdated and, in our opinion, an accurate identification of the species is not possible until a modern redescription is available (see Discussion for more details). Therefore, we identified our population as *M. cf. recens*. Nevertheless, given that our population fits the description of *M. recens*, it has to represent at least a closely related species and, as such, it can be used to estimate the phylogenetic position of *M. recens* s.str.

Genotyping

The DNA was extracted from individual animals following the Chelex[®] 100 resin (Bio-Rad) extraction method of Casquet *et al.* (2012) with modifications described in detail in Stec *et al.* (2015). We sequenced four DNA fragments, three nuclear (18S rRNA, 28S rRNA, ITS-2) and one mitochondrial (COI). All fragments were amplified and sequenced according to the protocols described in Stec *et al.* (2015); primers and original references for specific PCR programs are listed in Table 2. Sequencing products were read with the ABI 3130xl sequencer at the Molecular Ecology Lab, Institute of Environmental Sciences of the Jagiellonian University, Kraków, Poland. Sequences were processed in BioEdit v. 7.2.5 (Hall 1999) and submitted to GenBank.

Comparative molecular analysis

For molecular comparisons, all published sequences of the four above-mentioned markers for species of the *hufelandi* complex were downloaded from GenBank (listed in Table 3). Additionally, we also sequenced the four DNA fragments for a new population of *Macrobiotus macrocalix* Bertolani & Rebecchi, 1993 collected in Łękawica, southern Poland by DS in October 2015 (see Table 1 for sample details). The sequences were aligned using the default settings (in the case of COI) and the Q-INS-I method (in the case of ribosomal markers: 18S rRNA, 28S rRNA and ITS-2) of MAFFT v. 7 (Katoh *et al.* 2002; Katoh & Toh 2008) and manually checked against non-conservative alignments in BioEdit. Then, the aligned sequences were trimmed to 763 (18S rRNA), 712 (28S rRNA), 317 (ITS-2) and 618 (COI), bp. All COI sequences were translated into protein sequences in MEGA7 v. 7.0 (Kumar *et al.* 2016) to check against pseudogenes. According to the recommendation by Srivathsan & Meier (2012), uncorrected pairwise distances were calculated using MEGA7 instead of genetic distances corrected by Kimura 2 parameter model (K2P).

Data deposition

Raw morphometric measurements underlying the description of *Macrobiotus canaricus* sp. nov. are given in [Supplementary Materials](#) (SM.1) and are deposited in the Tardigrada Register (Michalczyk & Kaczmarek 2013) under www.tardigrada.net/register, whereas raw morphometric data underlying the description of the population of *M. cf. recens* are given in [Supplementary Materials](#) (SM.2). The DNA

STEC D. *et al.*, Tardigrades from the Canary Islands**Table 3.** Sequences of species of the *Macrobiotus hufelandi* group used for molecular comparisons and the phylogenetic analyses of *Macrobiotus canaricus* sp. nov. and *M. cf. recens* Cuénot, 1932. Underlined GenBank accession numbers indicate type or neotype sequences, whereas bolded numbers indicate new sequences obtained in this study.

DNA marker	Species	Accession number	Source
18S rRNA	" <i>M. hufelandi</i> " Schultze, 1834	GQ849024	Giribet <i>et al.</i> (1996)
	<i>M. hufelandi</i> group species	HQ604971, FJ435738–40	Bertolani <i>et al.</i> (2014), Guil & Giribet (2012)
	<i>M. hanna</i> e Nowak & Stec, 2018	<u>MH063922</u>	Nowak & Stec (2018)
	" <i>M. joannae</i> " Pilato & Binda, 1983	HQ604974–5	Bertolani <i>et al.</i> (2014)
	[= <i>M. hanna</i> e Nowak & Stec, 2018]		
	<i>M. kristenseni</i> Guidetti <i>et al.</i> , 2013	<u>KC193577</u>	Guidetti <i>et al.</i> (2013)
	<i>M. macrocalix</i> Bertolani & Rebecchi, 1993	<u>HQ604976</u> , MH063926	Bertolani <i>et al.</i> (2014), this study
	<i>M. papei</i> Stec <i>et al.</i> , 2018	<u>MH063881</u>	Stec <i>et al.</i> (2018d)
	<i>M. paulinae</i> Stec <i>et al.</i> , 2015	<u>KT935502</u>	Stec <i>et al.</i> (2015)
	<i>M. polypiformis</i> Roszkowska <i>et al.</i> , 2017	<u>KX810008</u>	Roszkowska <i>et al.</i> (2017)
	<i>M. polonicus</i> Pilato <i>et al.</i> , 2003	HM187580	Welnicz <i>et al.</i> (2011)
	<i>M. sapiens</i> Binda & Pilato, 1984	DQ839601	Bertolani <i>et al.</i> (2014)
28S rRNA	<i>M. scoticus</i> Stec <i>et al.</i> , 2017	<u>KY797265</u>	Stec <i>et al.</i> (2017b)
	<i>M. shonaicus</i> Stec <i>et al.</i> , 2018	<u>MG757132</u>	Stec <i>et al.</i> (2018a)
	<i>M. hanna</i> e Nowak & Stec, 2018	<u>MH063924</u>	Nowak & Stec (2018)
	<i>M. hufelandi</i> group species	FJ435751, FJ435754–5	Guil & Giribet (2012)
	<i>M. macrocalix</i> Bertolani & Rebecchi, 1993	MH063935	this study
	<i>M. papei</i> Stec <i>et al.</i> , 2018	<u>MH063880</u>	Stec <i>et al.</i> (2018d)
	<i>M. paulinae</i> Stec <i>et al.</i> , 2015	<u>KT935501</u>	Stec <i>et al.</i> (2015)
	<i>M. polypiformis</i> Roszkowska <i>et al.</i> , 2017	<u>KX810009</u>	Roszkowska <i>et al.</i> (2017)
	<i>M. scoticus</i> Stec <i>et al.</i> , 2017	<u>KY797266</u>	Stec <i>et al.</i> (2017b)
	<i>M. shonaicus</i> Stec <i>et al.</i> , 2018	<u>MG757133</u>	Stec <i>et al.</i> (2018a)
	<i>M. hanna</i> e Nowak & Stec, 2018	<u>MH063923</u>	Nowak & Stec (2018)
	<i>M. macrocalix</i> Bertolani & Rebecchi, 1993	MH063931	this study
ITS-2	<i>M. papei</i> Stec <i>et al.</i> , 2018	<u>MH063921</u>	Stec <i>et al.</i> (2018d)
	<i>M. paulinae</i> Stec <i>et al.</i> , 2015	<u>KT935500</u>	Stec <i>et al.</i> (2015)
	<i>M. polonicus</i> Pilato <i>et al.</i> , 2003	HM150647	Welnicz <i>et al.</i> (2011)
	<i>M. polypiformis</i> Roszkowska <i>et al.</i> , 2017	<u>KX810010</u>	Roszkowska <i>et al.</i> (2017)
	<i>M. sapiens</i> Binda & Pilato, 1984	GQ403680	Schill <i>et al.</i> (2010)
	<i>M. scoticus</i> Stec <i>et al.</i> , 2017	<u>KY797268</u>	Stec <i>et al.</i> (2017b)
	<i>M. shonaicus</i> Stec <i>et al.</i> , 2018	<u>MG757134–5</u>	Stec <i>et al.</i> (2018a)
	<i>M. hanna</i> e Nowak & Stec, 2018	<u>MH057764</u>	Nowak & Stec (2018)
COI	<i>M.cf. hufelandi</i> , Schultze, 1834	HQ876589–94, HQ876596	Bertolani <i>et al.</i> (2011a)
	<i>M. hufelandi</i> , Schultze, 1834	<u>HQ876584</u> , <u>HQ876586–8</u>	Bertolani <i>et al.</i> (2011a)
	<i>M. kristenseni</i> Guidetti <i>et al.</i> , 2013	<u>KC193575–6</u>	Guidetti <i>et al.</i> (2013)
	<i>M. macrocalix</i> Bertolani & Rebecchi, 1993	<u>FJ176203–07</u> , <u>FJ176208–17</u> , HQ876571, MH057767	Cesari <i>et al.</i> (2009), Bertolani <i>et al.</i> (2011a) this study
	<i>M. papei</i> Stec <i>et al.</i> , 2018	<u>MH057763</u>	Stec <i>et al.</i> (2018d)
	<i>M. paulinae</i> Stec <i>et al.</i> , 2015	<u>KT951668</u>	Stec <i>et al.</i> (2015)
	<i>M. polypiformis</i> Roszkowska <i>et al.</i> , 2017	<u>KX810011–2</u>	Roszkowska <i>et al.</i> (2017)
	<i>M. sandrae</i> Bertolani & Rebecchi, 1993	HQ876566–67, HQ876569–70, HQ876562–73, <u>HQ876574–83</u>	Bertolani <i>et al.</i> (2011a)
	<i>M. scoticus</i> Stec <i>et al.</i> , 2017	<u>KY797267</u>	Stec <i>et al.</i> (2017b)
	<i>M. shonaicus</i> Stec <i>et al.</i> , 2018	<u>MG757136–7</u>	Stec <i>et al.</i> (2018a)
	<i>M. terminalis</i> Bertolani & Rebecchi, 1993	JN673960, <u>AY598775</u>	Cesari <i>et al.</i> (2011), Guidetti <i>et al.</i> (2005)
	<i>M. vladimiri</i> Bertolani <i>et al.</i> , 2011	<u>HM136931–2</u> , <u>HM136933–4</u> , <u>HQ876568</u>	Bertolani <i>et al.</i> (2011a, 2011b)

sequences for the type population are deposited in GenBank (<https://www.ncbi.nlm.nih.gov/genbank>). Uncorrected pairwise distances are given in [Supplementary Materials](#) (SM.3).

Phylogenetic analysis

To verify the phylogenetic positions of the new species and *M. cf. recens*, phylogenetic trees were constructed using (1) all macrobiotid 18S rRNA sequences available from GenBank, (2) concatenated 18S rRNA+28S rRNA+ITS-2+COI sequences of macrobiotid species for which at least three of these markers were sequenced, and (3) all published *M. hufelandi* group COI sequences. In addition to the sequences of the *hufelandi* group listed in Table 3, we also used sequences of other species of the family Macrobiotidae published so far: Sands *et al.* (2008), Guidetti *et al.* (2009), Guil & Giribet (2012), Bertolani *et al.* (2014), Mapalo *et al.* (2016), Vecchi *et al.* (2016), Zawierucha *et al.* (2016), Mapalo *et al.* (2017), Stec & Kristensen (2017), Stec *et al.* (2018b). The sequences of *Milnesium variegidum* Morek *et al.*, 2016a and *Mi. berladnicorum* Ciobanu *et al.*, 2014 from Morek *et al.* (2016a) were used as the outgroup in the first two analyses, whereas four species of *Mesobiotus* served as the outgroup in the third analysis: *Me. ethiopicus* Stec & Kristensen, 2017, *Me. hilariae* Vecchi *et al.*, 2016, *Me. insanis* Mapalo *et al.*, 2017 and *Me. philippinicus* Mapalo *et al.*, 2016.

The sequences were aligned using the default settings (in the case of COI) and with the Q-INS-I method (in the case of ribosomal markers: 18S rRNA, 28S rRNA and ITS-2) of MAFFT v. 7 (Kato *et al.* 2002; Kato & Toh 2008) and then edited and checked manually in BioEdit. The alignments of 18S rRNA and COI sequences were trimmed to 739 bp and 618 bp, respectively, whereas the aligned sequences that were used to construct the concatenated data matrix were trimmed to: 728 bp (18S rRNA), 754 bp (28S rRNA), 570 bp (ITS-2) and 621 bp (COI). The sequences were concatenated in SequenceMatrix (Vaidya *et al.* 2011). The concatenated data matrix comprises species for which at least three of the aforementioned molecular markers are available. This resulted in only two gaps within the matrix: for the 18S rRNA sequence of *Me. insanis* (sequence too short) and the ITS-2 sequence of *Me. ethiopicus* (sequence not available). Using PartitionFinder v. 2.1.1 (Lanfear *et al.* 2016) under the Bayesian Information Criterion (BIC), the best scheme of partitioning and substitution models were chosen for posterior phylogenetic analysis. We ran the analysis to test all possible models implemented in the program. As the COI is a protein coding gene, before partitioning, we divided our alignments of this marker into 3 data blocks constituting separated three codon positions. As best-fit partitioning scheme, PartitionFinder always suggested the retention of all predefined partitions separately. See Table 4 for specific substitution models suggested for all tested data sets and partitions.

Bayesian inference (BI) marginal posterior probabilities were calculated for both data sets using MrBayes v. 3.2 (Ronquist & Huelsenbeck 2003). Random starting trees were used and the analysis was run for ten million generations, sampling the Markov chain every 1000 generations. An average standard deviation of split frequencies of < 0.01 was used as a guide to ensure the two independent analyses had converged. The program Tracer v. 1.7 (Rambaut *et al.* 2018) was then used to ensure Markov chains had reached stationarity and to determine the correct ‘burn-in’ for the analysis, which was the first 10% of generations. The ESS values were greater than 200 and a consensus tree was obtained after summarising the resulting topologies and discarding the ‘burn-in’. All final consensus trees were viewed and visualised by FigTree v. 1.4.3, available from <http://tree.bio.ed.ac.uk/software/figtree>.

Abbreviations

PCM = phase contrast light microscopy

SEM = scanning electron microscopy

IZiBB = Institute of Zoology and Biomedical Research, Jagiellonian University, Kraków, Poland

Table 4. The best scheme of partitioning and substitution models chosen for posterior phylogenetic analysis using PartitionFinder v. 2.1.1 under the Bayesian Information Criterion (BIC). The analyses were run to test all possible models implemented in the program.

Data set	Substitution model for the given partition					
	18S	28S	ITS-2	COI-1 st	COI-2 nd	COI-3 rd
18S	SYM+I+G	–	–	–	–	–
COI	–	–	–	SYM+I+G	GTR+I+G	TRN+G
18S+28S+ ITS-2+COI	SYM+G	GTR+G	SYM+G	GTR+I+G	GTR+I+G	GTR+G

Results

Taxonomic account

Phylum Tardigrada Doyère, 1840
 Class Eutardigrada Richters, 1926
 Order Parachela Schuster *et al.*, 1980
 Superfamily Macrobiotidea Thulin, 1928 (in Marley *et al.* 2011)
 Family Macrobiotidae Thulin, 1928
 Genus *Macrobiotus* C.A.S. Schultze, 1834

Macrobiotus canaricus sp. nov.

urn:lsid:zoobank.org:act:AE3AAEA9-D20E-4917-8ECD-12B30DB514B6

Figs 1–7, Tables 5–6

Etymology

The specific epithet refers to the Canary Islands, the place where the new species was found.

Material examined (162 animals, 57 eggs)

Specimens mounted on microscope slides in Hoyer's medium (120 animals + 42 eggs), fixed on SEM stubs (20 + 15), processed for DNA sequencing (7 animals), and aceto-orcein staining (15 animals).

Holotype

SPAIN: Canary Islands, Gran Canaria, Fagajesto, 28°03'05" N, 15°38'21" W, moss on a tree trunk in a pine forest (slide IZiBB ES.004.04).

Paratypes

SPAIN: 127 specimens, same data as for holotype (slides IZiBB ES.004.01–24); 42 eggs, same data as for holotype (slides IZiBB ES.004.25–33).

Description

Animals (measurements and statistics in Table 5)

Body white in adults, after fixation in Hoyer's medium transparent (Fig. 1A). Eyes present both in live animals and in specimens mounted in Hoyer's medium. Round and oval pores (0.4–0.7 µm in diameter), visible under PCM and SEM, scattered randomly on entire body cuticle (Fig. 2A–F), including external and internal surface of all legs (Fig. 2A–F). Extremely fine body granulation (granules 0.06–0.09 µm

Table 5. Measurements (in μm) of selected morphological structures of individuals of *Macrobiotus canaricus* sp. nov. mounted in Hoyer's medium (N = number of specimens/structures measured; Range = the smallest and the largest structure among all measured specimens; SD = standard deviation)

Character	N	Range		Mean		SD		Holotype	
		μm	pt	μm	pt	μm	pt	μm	pt
Body length	30	306–431	902–1206	368	1094	30	72	359	1088
Buccopharyngeal tube									
Buccal tube length	30	29.5–37.6	–	33.7	–	1.8	–	33.0	–
Stylet support insertion point	30	22.4–28.1	74.3–76.8	25.4	75.4	1.3	0.7	24.6	74.5
Buccal tube external width	30	3.2–5.4	10.3–15.5	4.3	12.8	0.4	1.1	4.3	13.0
Buccal tube internal width	30	2.1–3.3	6.6–9.5	2.7	7.9	0.3	0.8	2.5	7.6
Ventral lamina length	28	16.2–24.4	53.4–68.2	20.5	60.8	1.5	3.7	19.5	59.1
Placoid lengths									
Macroplacoid 1	30	5.5–9.5	17.7–28.4	7.3	21.5	1.0	2.4	8.0	24.2
Macroplacoid 2	30	4.6–7.1	14.6–20.5	5.8	17.2	0.6	1.4	6.0	18.2
Microplacoid	30	1.6–2.7	4.7–7.8	2.0	6.0	0.3	0.8	2.1	6.4
Macroplacoid row	30	11.5–17.5	36.8–52.0	14.1	41.8	1.4	3.1	14.4	43.6
Placoid row	30	14.0–19.4	42.6–55.3	16.6	49.2	1.5	3.0	17.0	51.5
Claw 1 lengths									
External primary branch	30	8.3–11.3	25.1–33.5	9.5	28.3	0.7	2.0	8.3	25.2
External secondary branch	30	6.2–9.3	19.0–26.2	7.4	22.1	0.7	1.9	6.5	19.7
Internal primary branch	29	7.5–10.6	23.8–32.0	9.2	27.4	0.6	1.9	8.6	26.1
Internal secondary branch	29	6.0–8.4	17.4–23.8	7.0	20.7	0.6	1.7	6.6	20.0
Claw 2 lengths									
External primary branch	30	8.6–11.8	25.3–35.6	9.8	29.3	0.7	2.2	9.0	27.3
External secondary branch	30	5.8–8.7	18.4–26.0	7.4	22.1	0.7	2.2	6.3	19.1
Internal primary branch	29	8.2–10.8	24.2–31.2	9.4	28.0	0.6	1.8	8.2	24.8
Internal secondary branch	29	5.4–8.6	15.9–25.1	7.2	21.5	0.6	2.0	7.4	22.4
Claw 3 lengths									
External primary branch	28	8.9–11.0	25.8–32.6	10.0	29.8	0.6	1.9	9.2	27.9
External secondary branch	28	6.4–9.7	18.8–28.4	7.7	22.8	0.7	2.4	7.2	21.8
Internal primary branch	28	5.6–11.3	16.5–31.6	9.5	28.1	1.0	3.0	8.1	24.5
Internal secondary branch	28	6.0–8.9	16.8–25.7	7.3	21.7	0.7	1.9	6.9	20.9
Claw 4 lengths									
Anterior primary branch	27	8.6–20.5	28.5–57.1	11.2	33.1	2.0	5.2	10.3	31.2
Anterior secondary branch	25	6.7–9.7	18.5–28.8	8.1	24.1	0.8	2.3	6.9	20.9
Posterior primary branch	26	9.6–13.2	28.3–37.6	11.2	33.2	0.8	2.7	10.5	31.8
Posterior secondary branch	23	6.6–9.5	19.4–27.3	8.1	24.0	0.9	2.3	7.4	22.4

in diameter), visible only under SEM, present on the dorso-posterior cuticle (Fig. 2E–F). Granulation patches on external surface of legs I–III clearly visible both under PCM and SEM (Fig. 3A–B). Granulation patches on internal surface of legs I–III weakly visible under PCM but clearly visible under SEM (Fig. 3C–D, empty indented arrowheads). Single, large, oval pore present at centre of each external patch on legs I–III (Fig. 3A–B, filled flat arrowheads). Cuticular bulge, resembling pulvinus, present on internal surface of legs I–III (Fig 3C–D, filled indented arrowheads). This structure is visible only if legs are fully extended and correctly oriented on slide. Cuticular granulation on legs IV present and always clearly visible both under PCM and SEM (Fig. 3E–F).

Mouth antero-ventral. Bucco-pharyngeal apparatus of the *Macrobiotus* type, with the ventral lamina and ten small peribuccal lamellae followed by six buccal sensory lobes (Fig. 4A–C). Under PCM, the oral cavity armature is of the *maculatus* type, i.e., only the third band of teeth is visible (Fig. 4A).

Under SEM, the oral cavity is always composed of three bands of teeth (Fig. 4B–C). The first band of teeth is composed of numerous extremely small cones arranged in one or two rows, situated anteriorly in the oral cavity, on the basal part of the peribuccal lamellae (Fig. 4B–C, filled arrowhead). The second band of teeth is situated between the ring fold and the third band of teeth and consists of cones, clearly larger than those of the first band (Fig. 4B–C, empty arrowhead). The teeth of the third band are located within the posterior portion of the oral cavity, between the second band of teeth and the buccal tube opening (Fig. 4B–C). The third band of teeth is discontinuous and divided into a dorsal and a ventral portion. Under PCM, the dorsal teeth form a transversal ridge weakly divided into three teeth, whereas the ventral teeth appear as two separate lateral transverse ridges between which a roundish median tooth is visible (Fig. 4A). Under SEM, the dorsal teeth are divided into three separate teeth: one median and two lateral, the median tooth has a slightly serrated edge (Fig. 4B). The ventral teeth are also separated into one median and two lateral teeth (Fig. 4C). The medio-ventral tooth is much smaller than the medio-dorsal tooth (Fig. 4B–C). Pharyngeal bulb spherical, with triangular apophyses, two rod-shaped macroplacoids and a small microplacoid (Fig. 4A). The first and the second macroplacoids have a fine central and a subterminal constriction, respectively. The macroplacoid length sequence is $2 < 1$.

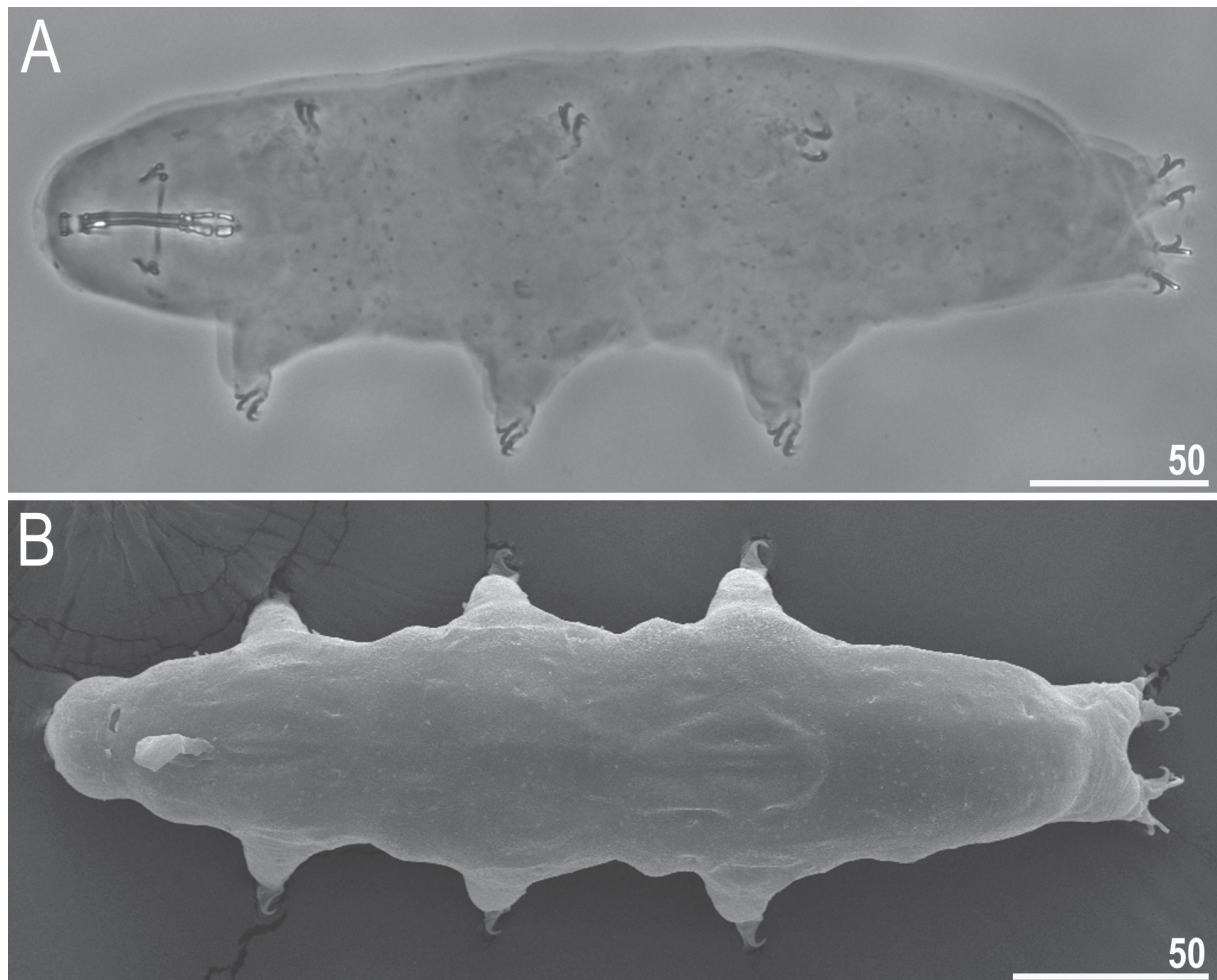


Fig. 1. *Macrobiotus canaricus* sp. nov., holotype, habitus. **A.** Dorso-ventral projection (Hoyer's medium, PCM). **B.** Dorsal view under SEM. Scale bars in μm .

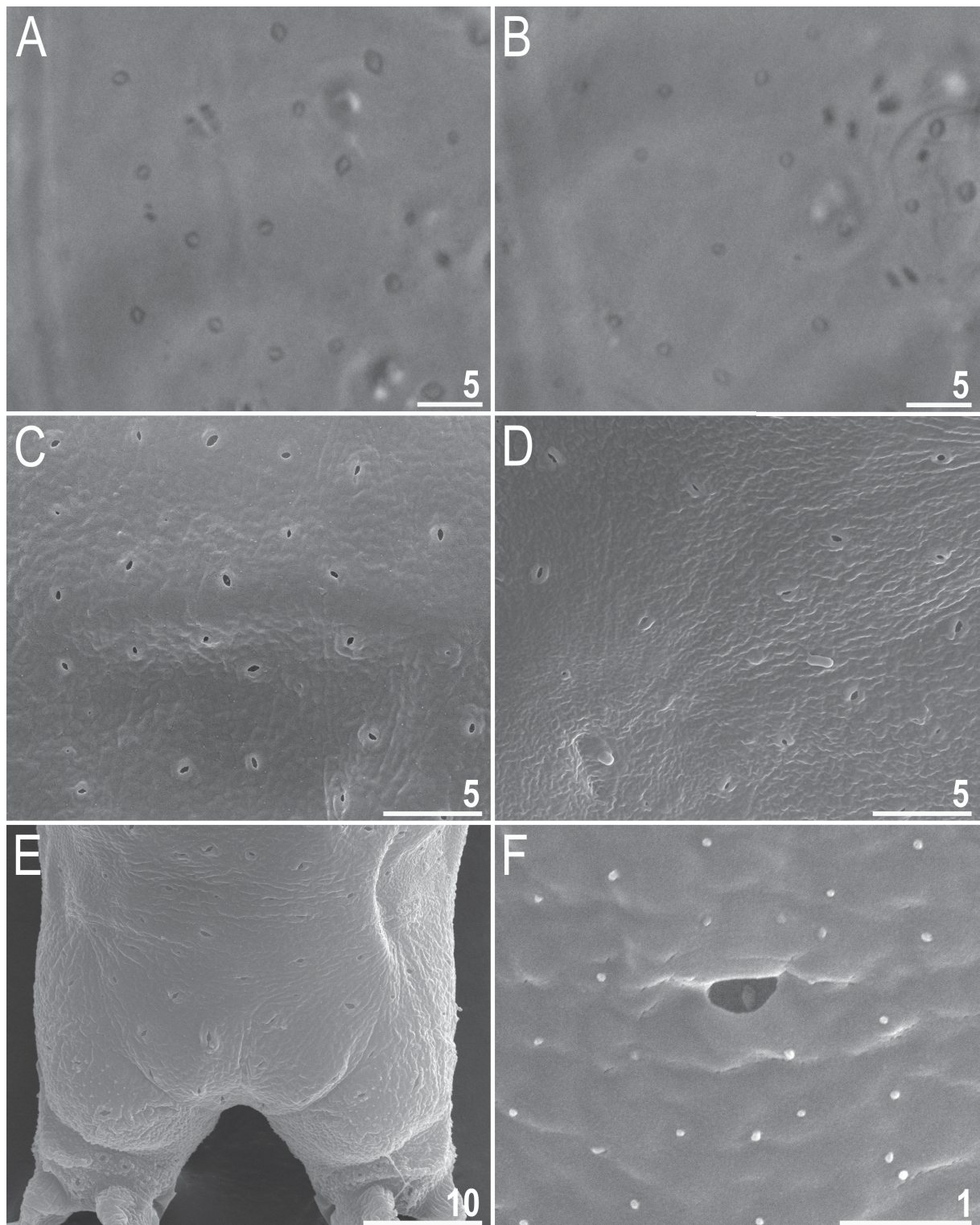


Fig. 2. *Macrobiotus canaricus* sp. nov., paratypes, body cuticle. **A–B.** Pores on the dorsal and ventral cuticle, respectively (PCM). **C–D.** Pores on the dorsal and ventral cuticle, respectively (SEM). **E.** Pores and fine granulation on the dorso-posterior cuticle (SEM). **F.** Close-up of a pore and granulation on the dorso-posterior cuticle (SEM). Scale bars in µm.

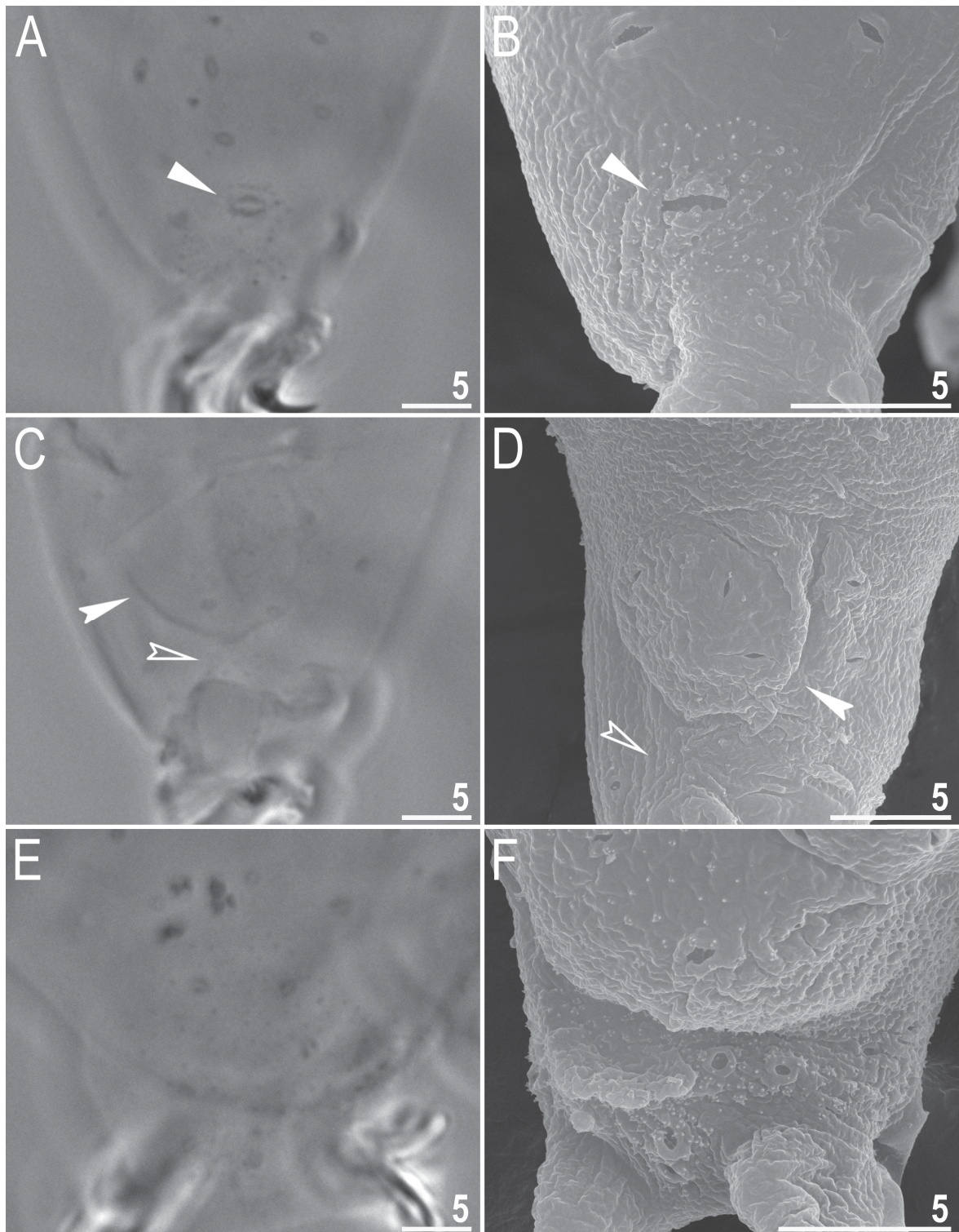


Fig. 3. *Macrobiotus canaricus* sp. nov., paratypes, cuticular structures on legs. **A–B.** The patch of granulation on the external surface of leg III with a large oval cuticular pore at the centre of the patch (PCM and SEM, respectively). **C–D.** The patch of granulation and the cuticular bulge (pulvinus) on the internal surface of leg II (PCM and SEM, respectively). **E–F.** Granulation on leg IV (PCM and SEM, respectively). Filled flat arrowheads indicate the large pore in the centre of the external granulation patch, filled indented arrowheads indicate the cuticular fold (pulvinus) on the internal leg surface, and empty indented arrowheads indicate granulation on the internal leg surface. Scale bars in µm.

Claws Y-shaped, of the *hufelandi* type (Fig. 5A–D). Primary branches with distinct accessory points and with an evident stalk connecting the claw to the lunula (Fig. 5A–D). Lunulae under all claws smooth (Fig. 5A–D). Cuticular bars under claws absent but muscle attachments are visible under claws I–III (Fig. 5A, C, filled arrowhead).

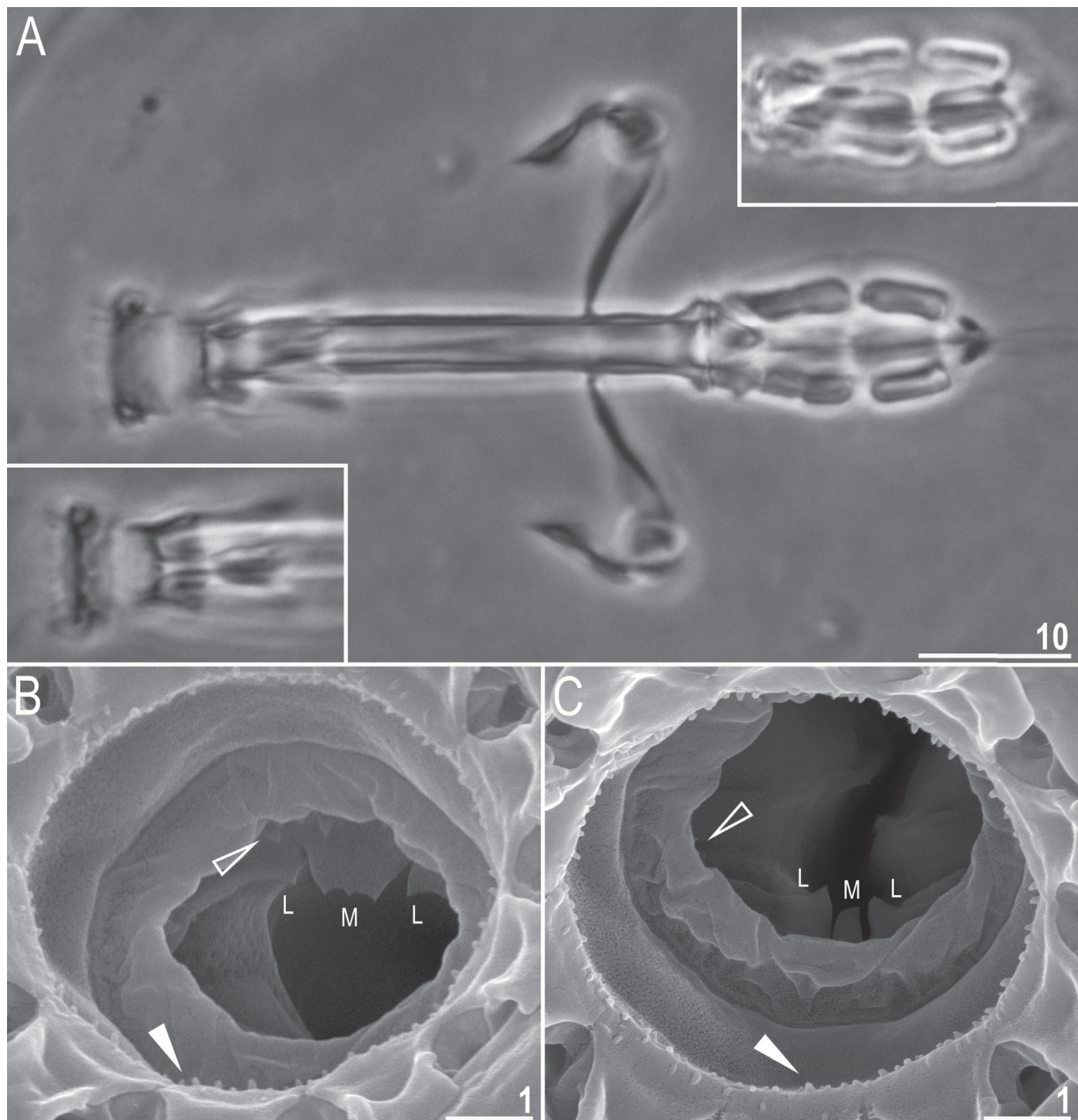


Fig. 4. *Macrobiotus canaricus* sp. nov., paratypes, buccal apparatus and the oral cavity armature seen under PCM and SEM. **A.** Dorsal projection of the entire buccal apparatus; the lower insert shows a ventral view of the oral cavity armature, the upper insert shows macroplacoid morphology (PCM). **B–C.** The oral cavity armature of a single paratype seen under SEM from different angles. Filled arrowheads indicate the teeth of the first band, empty arrowheads indicate the teeth of the second band, median teeth are marked with M, and lateral teeth are marked with L. Scale bars in μm .

Eggs (measurements and statistics in Table 6)

Laid freely, white, spherical or slightly oval (Figs 6A–B, 7A). The surface between processes of the *hufelandi* type, i.e., covered by a reticulum with very thin walls (Figs 6D–E, 7A–F). Peribasal meshes slightly larger and with slightly thicker walls compared to interbasal meshes (Figs 6D–E, 7B–F). The mesh diameter is always larger than mesh walls and nodes/knots (Figs 6D–E, 7B–F). The meshes are 0.3–1.0 µm in diameter, polygonal but with rounded edges. Under SEM, meshes deep and empty inside (Fig. 7C–F). Processes in the shape of inverted goblets with concave conical trunks and well-defined terminal discs (Figs 6C–F, 7A–F). Terminal discs strongly serrated, with a concave central area (Figs 6C–F, 7B–F). Sparse ultragranulation on the edges of terminal discs visible only under SEM (Fig. 7E–F). Three to five microgranules (0.25–0.30 µm in diameter), covered with ultragranulation, present in the centre of the terminal disc (visible only under SEM; Fig. 7B–F, empty arrowheads).

Reproductive mode

The examined population is dioecious (gonochoristic). Males were identified using aceto-orcein staining, which revealed testicles filled with spermatozoa. However, no morphological secondary sexual dimorphism, such as gibbosities on hind legs in males, was identified.

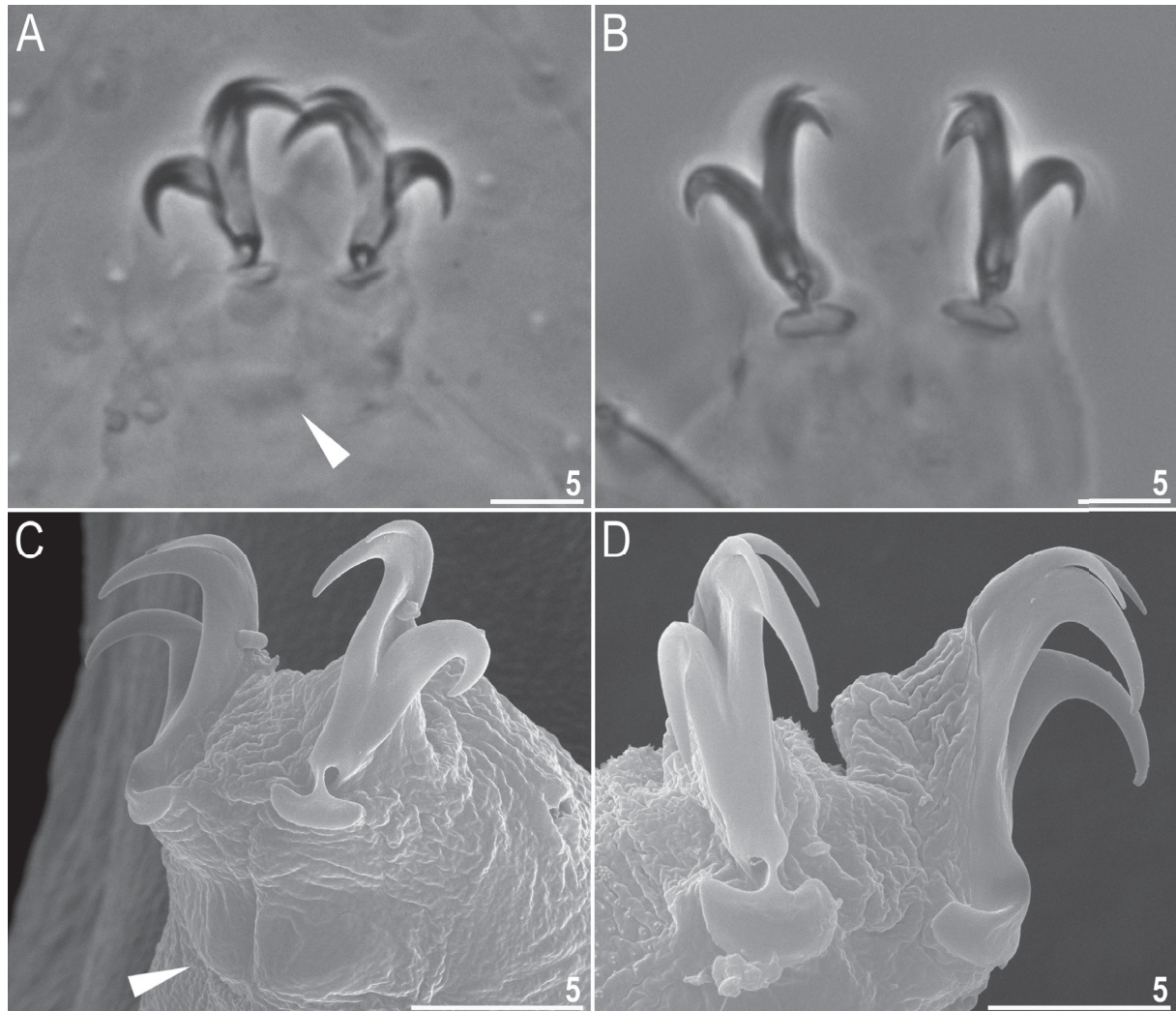


Fig. 5. *Macrobiotus canaricus* sp. nov., paratypes, claws. **A–B.** Claws III and IV seen under PCM. **C–D.** Claws I and IV seen under SEM. Arrowheads indicate faint muscle attachments under claws; Figs A and B assembled from several photos. Scale bars in µm.

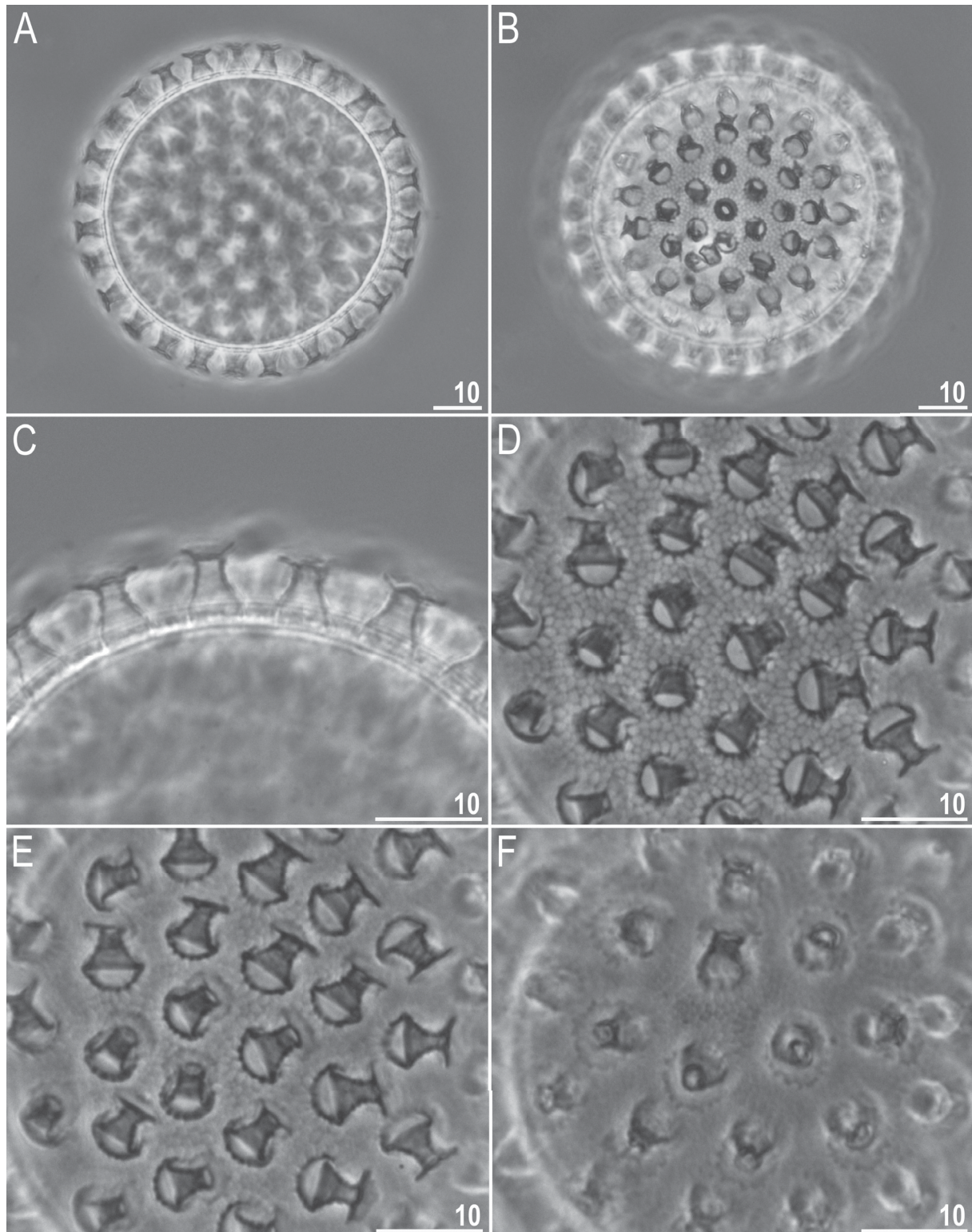


Fig. 6. *Macrobiotus canaricus* sp. nov., egg seen under PCM. **A.** Midsection under 400 × magnification. **B.** Surface under 400 × magnification. **C.** Midsection under 1000 × magnification. **D–E.** Surface under 1000 × magnification. **F.** Outlines of terminal discs of egg processes under 1000 × magnification. Scale bars in µm.

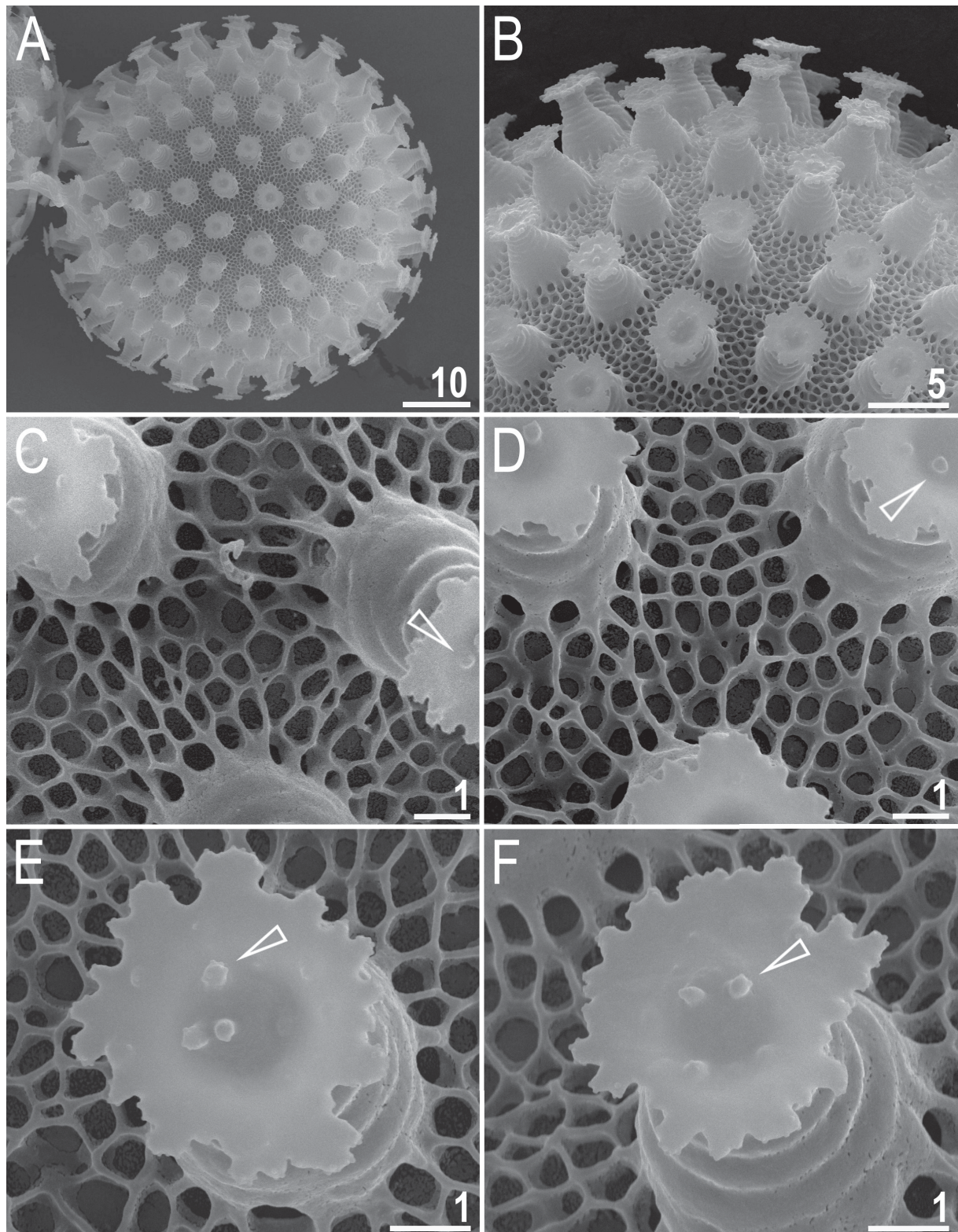


Fig. 7. *Macrobiotus canaricus* sp. nov., egg chorion morphology seen under SEM. **A.** Entire egg. **B.** Close up of the egg surface and processes. **C–D.** Details of the egg surface. **E–F.** Details of the terminal discs. Arrowheads indicate granules in the centre of the terminal discs. Scale bars in µm.

Table 6. Measurements (in μm) of selected morphological structures of the eggs of *Macrobiotus canaricus* sp. nov. mounted in Hoyer's medium. (N = number of eggs/structures measured; Range = the smallest and the largest structure among all measured specimens; SD = standard deviation)

Character	N	Range	Mean	SD
Egg bare diameter	30	59.9–81.6	68.7	5.0
Egg full diameter	30	70.2–96.2	80.8	5.4
Process height	90	3.9–8.2	5.5	1.0
Process base width	90	3.3–7.5	5.3	0.8
Process base/height ratio	90	60%–160%	99%	19%
Terminal disc width	90	3.4–7.5	4.8	0.8
Inter-process distance	90	1.5–5.5	3.4	0.8
Number of processes on the egg circumference	30	23–31	25.5	2.0

DNA sequences

We obtained sequences for all four of the above-mentioned molecular markers. The two conservative nuclear markers (18S rRNA, 28S rRNA) were represented by single haplotypes, whereas ITS-2 and COI exhibited three and two haplotypes, respectively. The p-genetic distance between the ITS-2 haplotypes ranged from 0.5 to 1.1% and between COI haplotypes it was equal to 1.3%. The 18S rRNA sequence (GenBank: MH063925) was 1033 bp long. The 28S rRNA sequence (GenBank: MH063934) was 721 bp long. The ITS-2 haplotypes 1–3 were 413 bp long (GenBank: MH063928, MH063929 and MH063930, respectively). The COI haplotypes 1–2 were 658 bp long (GenBank: MH057765 and MH057766, respectively).

Phenotypic differential diagnosis

By the oral cavity armature of the *maculatus* type and *hufelandi* type of egg shell ornamentation, smooth lunules under claws of all legs and granulation at least on legs IV, the new species is similar to *M. almadai* Fontoura *et al.*, 2008, *M. humilis* Binda & Pilato, 2001, and *M. rawsoni* Horning *et al.*, 1978, but can be differentiated specifically from:

Macrobiotus almadai, known only from the Azores (Fontoura *et al.* 2008), by the presence of the external and the internal patch of granulation on legs I–III (legs I–III smooth in *M. almadai*) and by the presence of a single large pore in the centre of the external patch on legs I–III (occasionally, regular cuticular pores may be present on some legs, but such pores are small and never present on all legs in the same place in *M. almadai*).

Macrobiotus humilis, reported only from its type locality in Sri Lanka (Binda & Pilato 2001), by the presence of three separated dorsal teeth of the third band (dorsal teeth fused into a single transversal ridge in *M. humilis*), the presence of a subterminal constriction in the second macroplacoid (second macroplacoid without constrictions in *M. humilis*), more posteriorly inserted stylet supports ($pt = 74.3$ – 76.8 in the new species vs $pt = 71.1$ – 71.3 in *M. humilis*), slightly higher pt of the second macroplacoid length ($pt = 14.6$ – 19.6 in the new species vs $pt = 12.5$ – 14.4 in *M. humilis*) and by irregularly serrated edges of the terminal discs on egg processes (edges of terminal discs regularly indented in *M. humilis*).

Macrobiotus rawsoni, known only from its type locality in New Zealand (Horning *et al.* 1978; Kaczmarek & Michalczyk 2017a), by the presence of granulation on all legs (granulation present only on legs IV in *M. rawsoni*), the presence of a subterminal constriction in the second macroplacoid (second macroplacoid without constrictions in *M. rawsoni*), the absence of cuticular bars under the claws on

legs I–III (thin paired bars present in *M. rawsoni*), more anteriorly inserted stylet supports ($pt = 74.3–76.8$ in the new species vs $pt = 77.0–77.1$ in *M. rawsoni*), a different morphology of reticulation on the egg surface between processes (several lines of mesh between neighbouring egg processes in the new species vs two lines of mesh between neighbouring egg processes in *M. rawsoni*) and by a smaller mesh size in the chorion reticulum ($0.3–1.0\ \mu\text{m}$ in diameter in the new species vs $1.8–2.5\ \mu\text{m}$ in diameter in *M. rawsoni*).

Genotypic differential diagnosis

The ranges of uncorrected genetic p-distances between the new species and species of the *Macrobotus hufelandi* complex, for which sequences are available from GenBank, are as follows:

- **18S rRNA:** 0.5–3.7% (2.0% on average), being most similar to two undetermined species of the *M. hufelandi* group from Spain (FJ435738–9) and to *M. macrocalix* from Poland (MH063926) and the least similar to *M. polypiformis* Roszkowska *et al.*, 2017 from Ecuador (KX810008)
- **28S rRNA:** 1.9–13.2% (6.1% on average), being most similar to three undetermined species of the *M. hufelandi* complex from Spain (FJ435751 and FJ435754–5) and the least similar to *M. polypiformis* from Ecuador (KX810009)
- **ITS-2:** 5.3–27.8% (17.0% on average), with the most similar being *M. macrocalix* from Poland (MH063931) and the least similar being *M. polypiformis* from Ecuador (KX810010)
- **COI:** 17.2–24.7% (19.2% on average), with the most similar being *M. hanna*e Nowak & Stec, 2018 from Poland (MH057764) and the least similar being *M. papei* Stec *et al.*, 2018 from Tanzania (MH057763)

Macrobotus cf. *recens* Cuénot, 1932

Figs 8–14, Tables 7–8

Material examined (114 animals, 48 eggs)

SPAIN: Canary Islands, Gran Canaria, Presa de Lugarejos, $28^{\circ}02'38''\text{ N}$, $15^{\circ}40'22''\text{ W}$, 885 m a.s.l., lichen on a stone wall. Specimens mounted on microscope slides in Hoyer's medium (87 animals + 38 eggs), fixed on SEM stubs (8+10), processed for DNA sequencing (4 animals) and used for aceto-orcein staining (15 animals). Slide depositories: 87 animals (slides: ES.006.*, where the asterisk can be substituted by the following numbers 1–4, 10–12, 14–17) and 38 eggs (slides: ES.006.*: 5–9, 13) (IZiBB).

Description of the population from Gran Canaria

Animals (measurements and statistics in Table 7)

Body white in juveniles and slightly yellowish in adults, after fixation in Hoyer's medium transparent (Fig. 8A). Eyes present in live animals and in specimens mounted in Hoyer's medium. Elliptical and sometimes roundish pores ($1.0–1.8\ \mu\text{m}$ in diameter), visible under PCM and SEM, scattered randomly on entire body cuticle (Fig. 8B–E), including the external and internal surface of all legs (Fig. 9A–I). Inside pores several granules, visible only under SEM, always present (Fig. 8E). Granulation patch on external and internal surfaces of legs I–III present (Fig. 9A–E). Single pore present at centre of each external granulation patch (Fig. 9A–C). Granulation patch on external surface larger and more distinct than the one on internal surface (Fig. 9A–E). Faint cuticular fold present on external surface of legs I–III just above claws (Fig. 9A–B, empty arrowhead), whereas on internal surface of legs I–III there is a cuticular bulge resembling pulvinus (Fig. 9D–E, filled arrowhead). Both external fold and internal bulge visible only if legs are fully extended and correctly oriented on slide (particularly cuticular fold above claws). Granulation on legs IV always clearly visible and consists of two granulation patches: the distal

Table 7. Measurements (in μm) of selected morphological structures of individuals of *Macrobiotus* cf. *recens* Cuénot, 1932 from the Canary Islands mounted in Hoyer's medium. (N = number of specimens/structures measured; Range = the smallest and the largest structure among all measured specimens; SD = standard deviation)

Character	N	Range		Mean		SD	
		μm	<i>pt</i>	μm	<i>pt</i>	μm	<i>pt</i>
Body length	30	353–731	684–1174	576	1042	66	90
Buccopharyngeal tube							
Buccal tube length	30	48.5–70.3	–	55.4	–	5.8	–
Stylet support insertion point	30	39.3–57.2	79.1–83.5	44.7	80.8	4.9	1.0
Buccal tube external width	30	8.0–15.6	15.4–22.8	9.8	17.6	1.6	1.4
Buccal tube internal width	30	6.1–12.2	11.6–17.8	7.5	13.4	1.3	1.3
Ventral lamina length	30	24.9–42.3	47.3–63.8	32.9	59.6	3.2	4.2
Placoid lengths							
Macroplacoid 1	30	9.4–19.4	18.1–30.5	13.3	24.0	2.3	2.6
Macroplacoid 2	30	8.2–17.0	15.9–24.8	10.6	19.0	2.0	1.8
Microplacoid	30	3.1–8.4	5.8–12.3	4.7	8.5	1.1	1.3
Macroplacoid row	30	12.5–39.1	24.1–57.1	25.4	45.7	5.1	6.2
Placoid row	30	17.3–52.6	33.4–76.8	32.7	58.7	6.1	6.3
Claw 1 lengths							
External primary branch	30	12.3–18.0	23.1–32.7	14.6	26.6	1.5	2.4
External secondary branch	29	9.6–14.6	17.9–23.8	11.7	21.1	1.2	1.6
Internal primary branch	30	12.0–17.0	22.7–29.1	14.1	25.6	1.2	1.9
Internal secondary branch	30	8.2–13.7	15.7–24.5	11.1	20.1	1.3	2.4
Claw 2 lengths							
External primary branch	30	12.8–18.3	23.2–31.6	15.0	27.2	1.3	2.3
External secondary branch	30	8.0–14.6	16.5–26.4	11.7	21.2	1.5	2.5
Internal primary branch	30	12.0–20.5	22.2–39.7	14.7	26.6	1.8	3.3
Internal secondary branch	30	8.5–14.9	15.5–25.6	11.2	20.4	1.6	2.7
Claw 3 lengths							
External primary branch	30	13.3–18.2	23.9–33.7	15.2	27.6	1.2	2.4
External secondary branch	30	8.9–16.2	17.2–26.1	11.7	21.2	1.5	2.0
Internal primary branch	30	11.1–17.5	20.4–32.7	14.5	26.3	1.4	2.5
Internal secondary branch	30	9.1–15.1	16.7–29.2	11.5	20.8	1.3	2.5
Claw 4 lengths							
Anterior primary branch	29	11.8–22.2	22.6–36.7	17.3	31.3	1.9	3.2
Anterior secondary branch	28	11.1–17.2	20.6–29.0	13.1	23.7	1.5	2.0
Posterior primary branch	27	14.5–23.4	28.0–38.9	18.1	32.7	1.6	2.5
Posterior secondary branch	26	11.1–17.4	20.2–32.0	13.7	24.6	1.6	3.1

patch with densely distributed granules situated just above claws and the proximal patch being wider with more sparsely distributed granules located immediately above distal patch (Fig. 9G–I).

Mouth antero-ventral. Bucco-pharyngeal apparatus of the *Macrobiotus* type (Fig. 10A–C), with ventral lamina and ten small peribuccal lamellae followed by six buccal sensory lobes. Under PCM, oral cavity armature of the *hufelandi* type, i.e., with all three bands of teeth always visible (Fig. 10B–C). First band of teeth composed of numerous very small cones arranged in four to six rows situated anteriorly in oral cavity, just behind bases of peribuccal lamellae (Figs 10B–C, 11A–B, filled arrowhead). Second band of teeth situated between ring fold and third band of teeth and comprises 4–5 rows of small cones, slightly larger than those of first band (Figs 10B–C, 11A–B, empty arrowhead). Teeth of the third band located within posterior portion of oral cavity, between the second band of teeth and buccal tube

opening (Figs 10B–C, 11A–B). Third band of teeth discontinuous and divided into dorsal and ventral portions. Under PCM, dorsal teeth seen as three distinct transversal ridges, whereas ventral teeth appear as two separate lateral transversal ridges and a roundish median tooth (Fig. 10B–C). Under SEM, both dorsal and ventral teeth clearly distinct (Fig. 11A–B). Medio-ventral tooth rarely divided into two or three smaller teeth (Fig. 11B). Under SEM, margins of dorsal and latero-ventral teeth slightly serrated (Fig. 11A–B). Pharyngeal bulb spherical, with triangular apophyses, two rod-shaped macroplacoids and

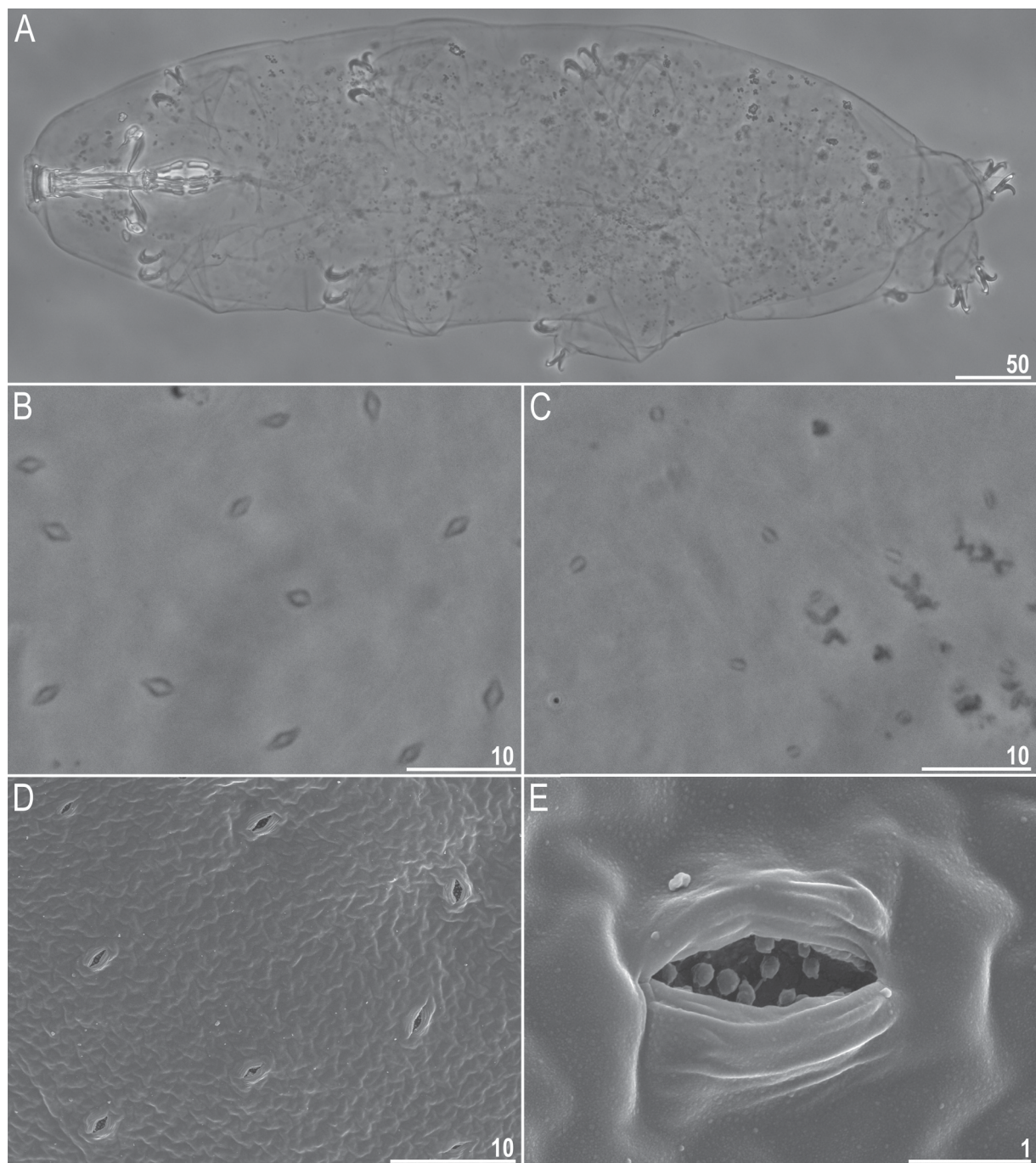


Fig. 8. *Macrobiotus* cf. *recens* Cuénnot, 1932, Gran Canaria, habitus and cuticular pores. **A.** Dorso-ventral projection (PCM). **B–C.** Pores on the dorsal and ventral cuticle, respectively (PCM). **D.** Pores on the dorsal cuticle (SEM). **E.** Granulation inside a single pore (SEM). Scale bars in µm.

one big rod-shaped microplacoid (Fig. 10D–E). Macroplacoid length sequence $2 < 1$. First and second macroplacoid with the central and the subterminal constriction, respectively (Fig. 10D–E).

Claws Y shaped, of the *hufelandi* type (Fig. 12A–D). Primary branches with distinct accessory points and with an evident stalk connecting claw to lunula (Fig. 12A–D). Lunulae I–III smooth (Fig. 12A, C), whereas lunulae IV slightly crenulated (Fig. 12B, D). Faint cuticular bars under claws I–III present, more visible in larger specimens (Fig. 12A, C). Horseshoe-shaped structure connects anterior and posterior lunules (Fig. 12B).

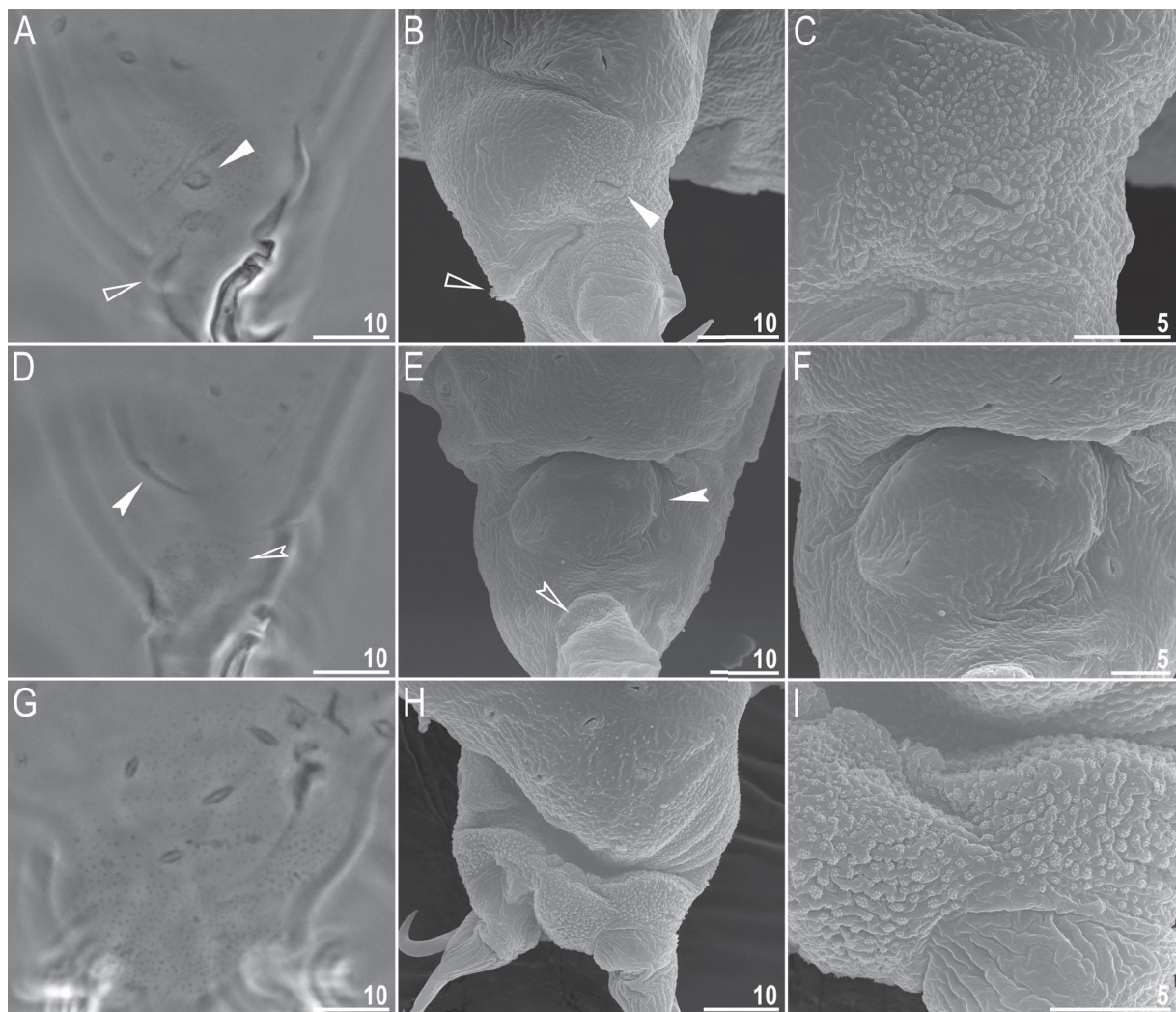


Fig. 9. *Macrobiotus cf. recens* Cuénot, 1932, Gran Canaria, structures on forelegs seen under PCM and under SEM. **A–C.** The patch of granulation on the external surface of leg II, with the cuticular pore at the centre of the patch and the cuticular fold above the claws. **D–F.** The patch of granulation and the cuticular bulge (pulvinus) on the internal surface of leg III. **G–I.** Granulation on leg IV. Filled flat arrowheads indicate the large pore at the centre of the external granulation patch, empty flat arrowheads indicate the cuticular fold above the claws on the external leg surface, filled indented arrowheads indicate the cuticular fold (pulvinus) on the internal leg surface, and empty indented arrowheads indicate granulation on the internal leg surface. Scale bars in µm.

Eggs (measurements and statistics in Table 8)

Laid freely, white/light yellow, spherical or slightly oval (Figs 13A–B, 14A). Surface between processes of the *hufelandi* type, i.e., chorion surface between processes covered by reticulum with very small meshes (Figs 13C–D, 14B–E). Under PCM, surface between processes seems to be covered by dark dots (Fig. 13C) and only sometimes a clear reticulation is visible (Fig. 13D). Several rows of meshes between egg processes (usually 7–8). Mesh borders and nodes/knots thick and sometimes wider than mesh diameter (Fig. 14B–E). Meshes circular or slightly oval (0.4–0.9 μm in diameter) and under SEM

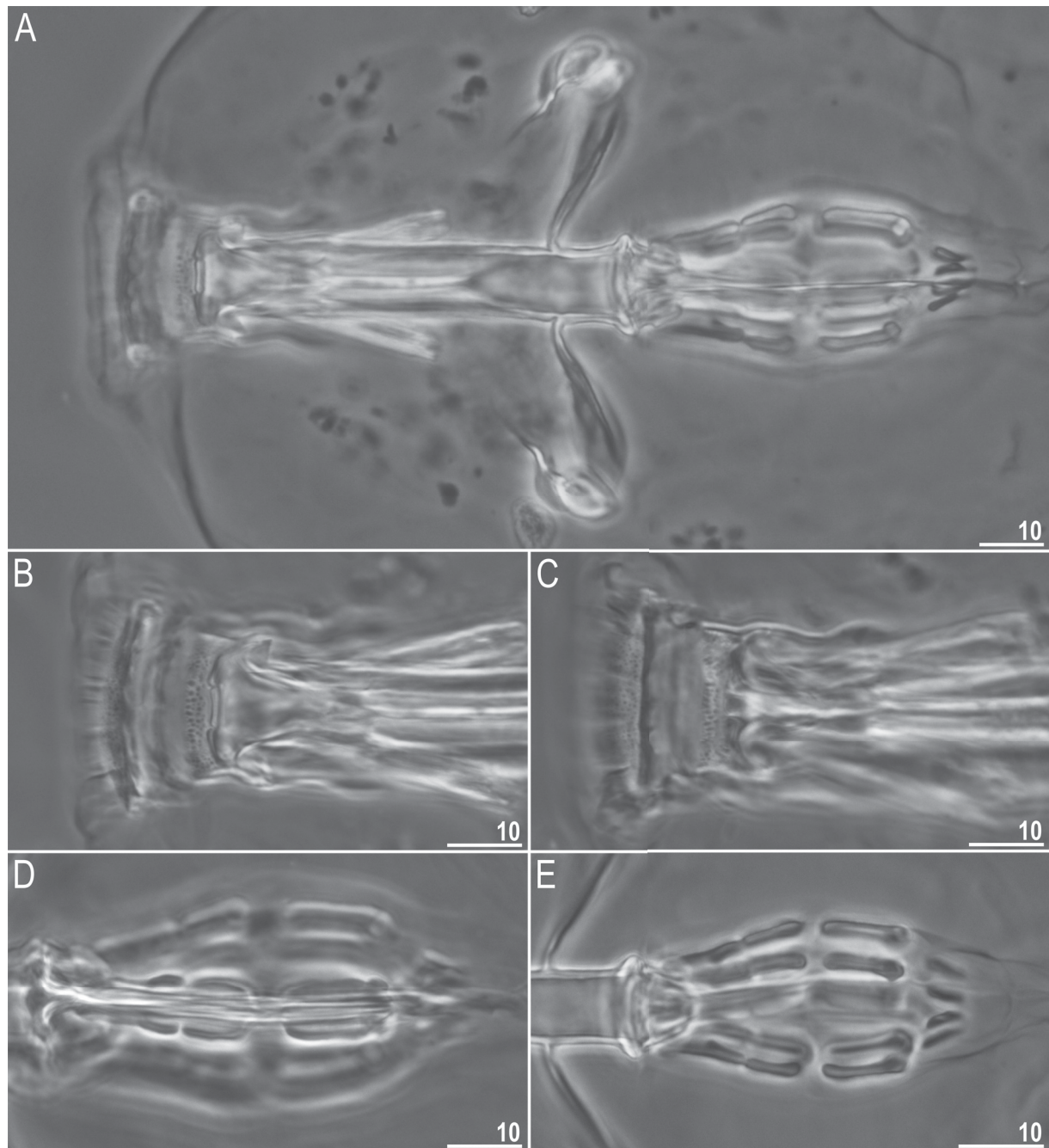


Fig. 10. *Macrobiotus cf. recens* Cuénot, 1932, Gran Canaria, buccal apparatus and the oral cavity armature seen under PCM. **A.** Dorsal projection of the entire buccal apparatus. **B–C.** Dorsal and ventral view of the oral cavity armature, respectively. **D–E.** Placoid morphology. Scale bars in μm .

Table 8. Measurements (in μm) of selected morphological structures of the eggs of *Macrobiotus* cf. *recens* Cuénot, 1932 from the Canary Islands mounted in Hoyer's medium. (N = number of eggs/structures measured; Range = the smallest and the largest structure among all measured specimens; SD = standard deviation)

Character	N	Range	Mean	SD
Egg bare diameter	30	86.1–113.7	102.1	8.0
Egg full diameter	30	113.6–151.4	137.6	9.3
Process height	90	13.4–26.3	20.0	2.7
Process base width	90	3.2–10.7	6.3	1.4
Process base/height ratio	90	16%–45%	32%	6%
Inter-process distance	90	3.7–74.0	6.6	7.3
Number of processes on the egg circumference	30	21–27	24.6	1.5

all pores empty inside (Fig. 14C–D). Mesh diameter decreases gradually from peribasal to interbasal meshes (Fig. 14C–D). Short thickenings radiating from process bases are often visible under PCM and always visible under SEM; thickenings may create a crown around process bases (Figs 13C, 14C–D, filled arrowheads). Processes in the shape of high and thin cones with straight conical trunks, devoid of terminal discs and sometimes bifurcated (Figs 13A–F, 14A–F). Trunks of processes slightly undulated. Undulations covered by numerous granules composed of microgranule aggregations. Undulations and granules poorly visible under PCM (Fig. 13C–F). Undulations, granules and microgranules always clearly visible under SEM (Fig. 14B, E–F).

Reproductive mode

The examined population is dioecious (gonochoristic). Males were identified using aceto-orcein staining, which revealed testicles filled with spermatozoa. However, no morphological secondary sexual dimorphism, such as gibbosities on hind legs in males, was identified.

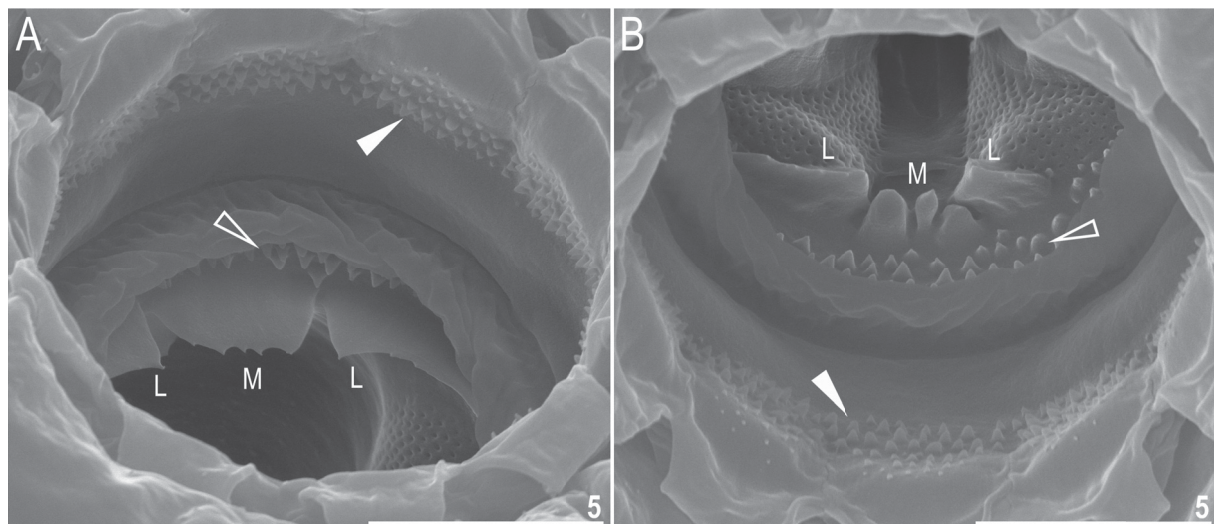


Fig. 11. *Macrobiotus* cf. *recens* Cuénot, 1932, Gran Canaria, oral cavity armature seen under SEM from different angles. **A.** View of the dorsal portion. **B.** View of the ventral portion. Filled arrowheads indicate the teeth of the first band, empty arrowheads indicate the teeth of the second band, median teeth are marked with M, and lateral teeth are marked with L. Scale bars in μm .

DNA sequences

We obtained sequences for all four of the above-mentioned molecular markers. The two conservative nuclear markers (18S rRNA, 28S rRNA) were represented by single haplotypes, whereas both ITS-2 and COI exhibited two haplotypes. The p-genetic distance between the ITS-2 as well as between the COI haplotypes was 1.1%. The 18S rRNA sequence (GenBank: MH063927) was 1033 bp long, the 28S rRNA sequence (GenBank: MH063936) was 725 bp long, the ITS-2 haplotype 1 and 2 sequences (GenBank: MH063932 and MH063933, respectively) were 420 bp long; the COI haplotype 1 and 2 sequences (GenBank: MH057768 and MH057769, respectively) were 658 bp long.

Genotypic differential diagnosis

The ranges of uncorrected genetic p-distances between the new species and species of the *Macrobiotus hufelandi* complex, for which sequences are available from GenBank, are as follows:



Fig. 12. *Macrobiotus cf. recens* Cuénot, 1932, Gran Canaria, claws. **A–B.** Claws III and IV seen under PCM, with smooth and weakly crenulated lunules, respectively. **C–D.** Claws II and IV seen under SEM, with smooth and weakly crenulated lunules, respectively. A and B assembled from several photos. Filled arrowheads indicate faint cuticular bars under claws whereas empty arrowhead indicates a horseshoe-shaped structure. Scale bars in μm.

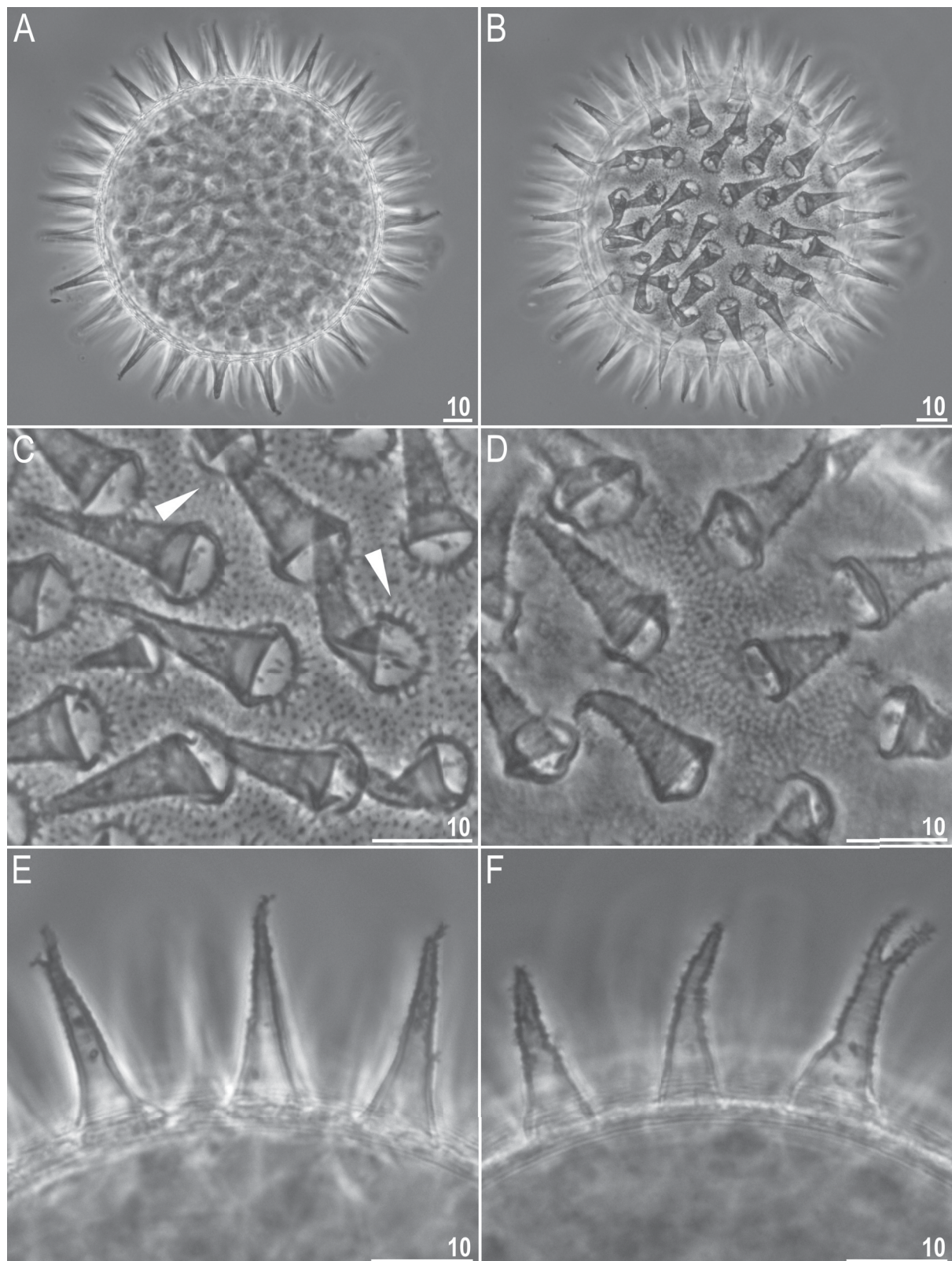


Fig. 13. *Macrobiotus cf. recens* Cuénot, 1932, Gran Canaria, egg seen under PCM. **A.** Midsection under $400\times$ magnification. **B.** Surface under $400\times$ magnification. **C–D.** Surface under $1000\times$ magnification. **E–F.** Midsection under $1000\times$ magnification. Filled arrowheads indicate short thickenings radiating from the processes bases. Scale bars in μm .

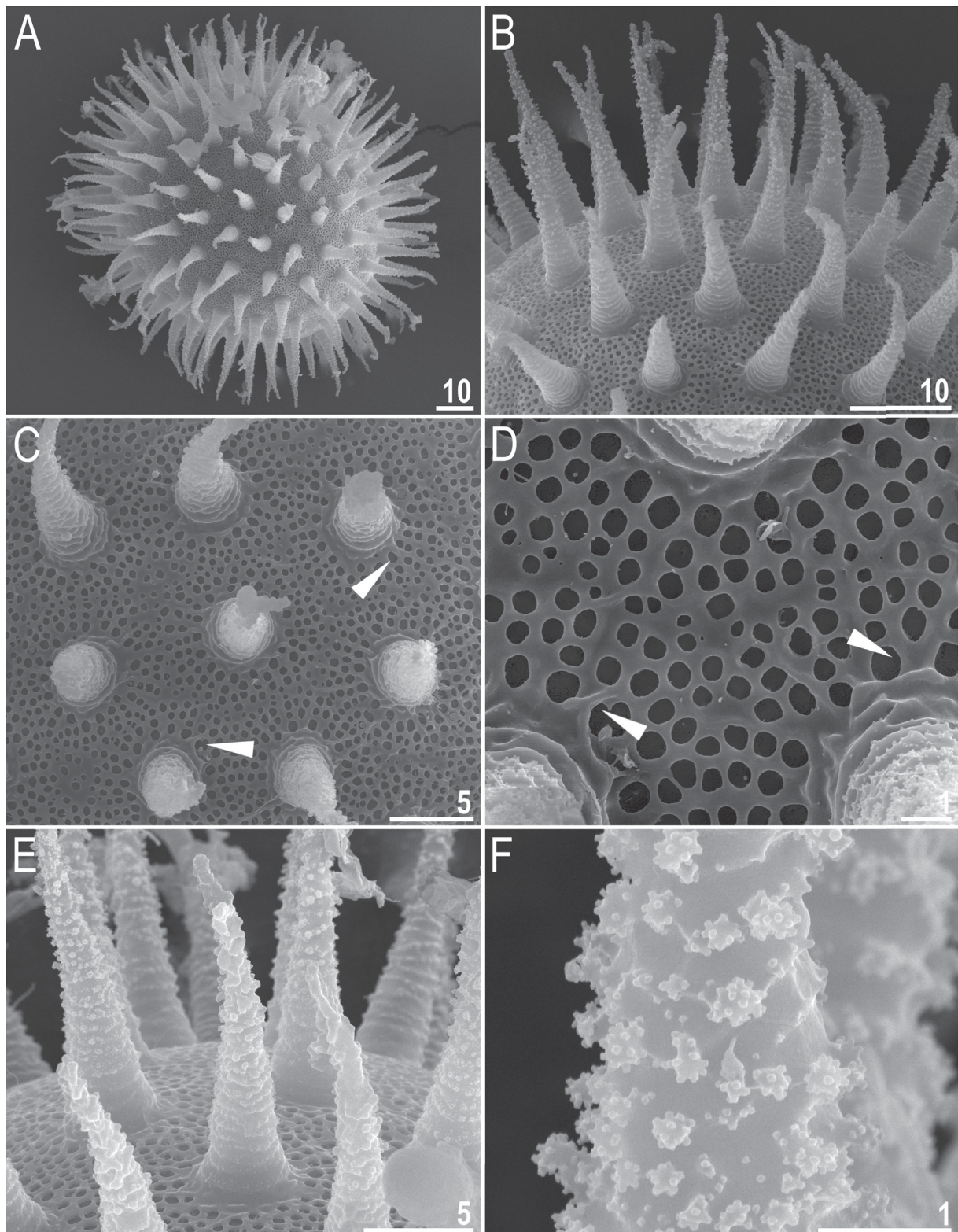


Fig. 14. *Macrobiotus cf. recens* Cuénot, 1932, Gran Canaria, egg chorion morphology seen under SEM. **A.** Entire egg. **B.** Close up of the egg surface and processes. **C–D.** Details of the egg surface. **E–F.** Details of process morphology. Filled arrowheads indicate short thickenings radiating from the process bases. Scale bars in µm.

- **18S rRNA:** 0.4–4.0% (2.0% on average), with the most similar being *M. macrocalix* from Poland (MH063926) and the least similar being *M. polypiformis* from Ecuador (KX810008)
- **28S rRNA:** 1.3–13.6% (6.5% on average), with the most similar being *M. macrocalix* from Poland (MH063935) and the least similar being *M. polypiformis* from Ecuador (KX810009)
- **ITS-2:** 3.7–26.7% (16.9% on average), with the most similar being *M. macrocalix* from Poland (MH063931) and the least similar being *M. polypiformis* from Ecuador (KX810010)
- **COI:** 16.4–24.7% (19.1% on average), with the most similar being *M. macrocalix* from Italy (FJ176203–7 and FJ176213–7) and the least similar being *M. papei* from Tanzania (MH057763)

Molecular phylogeny

Phylogenetic analyses conducted on macrobiotid 18S rRNA sequences as well as on the concatenated macrobiotid data set unambiguously confirmed that the two studied species represent the *M. hufelandi* group (Figs 15–16). The phylogeny based on the COI sequences of the *hufelandi* group also corroborated these results, since none of the two species were recovered external to the species of the *hufelandi* group (Fig. 17). In all analyses, two clades within the *hufelandi* group were present, although the species composition varied slightly between phylogenies based on different markers. One clade grouped exclusively species that exhibit modified egg processes (*M. paulinae* Stec *et al.*, 2015, *M. polypiformis*, *M. papei*, *M. shonaicus* Stec *et al.*, 2018a, *M. scoticus* Stec *et al.*, 2017b and *M. kristenseni* Guidetti, Peluffo, Rocha, Cesari & Moly de Peluffo, 2013); ‘the *kristenseni* clade’ henceforth. The other clade comprised mostly species with typical inverted goblet-shaped egg processes (‘the *hufelandi* clade’ hereafter). In contrast to our predictions, *M. cf. recens*, with its atypical egg processes, was always embedded within the *hufelandi* clade. The two clades were well supported in phylogenies based on the concatenated data set and on COI sequences, but weakly supported in the 18S rRNA tree (Figs 15–17). Moreover, in the 18S rRNA analysis the *kristenseni* clade, in addition to the majority of species with modified egg processes, comprised *M. sapiens* Binda & Pilato, 1984 (DQ839601) and undetermined species of the *M. hufelandi* group (HQ604971), of which at least the first species exhibits the typical egg morphology. In contrast to other analyses, the 18S rRNA phylogeny recovered a clade, with *X. pseudohufelandi* (Iharos, 1966) and *M. polonicus* Pilato, Kaczmarek, Michalczyk & Lisi, 2003, that was in a sister relationship to all other species of the *hufelandi* group, suggesting that the *hufelandi* group is polyphyletic or that *Xerobiotus* belongs to the *hufelandi* group (Fig. 15).

Discussion

Thanks to the detailed morphological and molecular analyses, we were able to describe a new species *M. canaricus* sp. nov. and characterise a population of *M. cf. recens*, both collected on the Canary Islands. Moreover, by the use of phylogenetic inference, we confirmed the affinity of both studied species with the *M. hufelandi* group.

Macrobiotus recens was described in 1932 by Lucien Cuénot, but soon after, some researchers started questioning the status of this species and classified it as ‘*M. hufelandi* forma *recens*’ (Marcus 1936; Ramazzotti 1945, 1962, 1972; Rudescu 1946; Grigarick *et al.* 1973). However, Maucci (1979) analysed several Portuguese populations he identified as *M. recens* morphologically and confirmed that the taxon is indeed a ‘bona species’. However, it should be noted that the locus typicus for *M. recens* is La Tardière, a commune in the Vendée department, in the Pays de la Loire region in western France. Thus, similarly to our study, the comparison presented by Maucci (1979) has to be treated with caution, as it may be based on a related species rather than *M. recens* s.str. Since the original description is outdated and incomplete, a confident identification of *M. recens* is currently not possible. Although Pilato & Bertolani (2004) supported the identification of the Portuguese populations studied by Maucci (1979) as *M. recens*, we think that without an integrative redescription of the species it is not possible to test whether *M. recens* is a single species or a complex of morphologically similar taxa. Thus, instead of designating the Canary

Islands population as *M. recens*, we treat it as *M. cf. recens*, at the same time highlighting the need for a redescription based on new material from the locus typicus. It could be hypothesised that our Canarian population, as well as previous records of *M. recens* outside the type locality, i.e., from Romania (Rudescu 1946), Italy (Ramazzotti 1945), Greece (Marcus 1936), Portugal (Maucci 1979) and China (Rahm 1937; Bartoš 1963), may represent distinct taxa that could become identifiable only after a modern redescription of *M. recens*. This would not be surprising, as such a pattern has been observed before, for example, when nominal taxa in other groups were described: *Macrobiotus hufelandi* (redescribed by Bertolani & Rebecchi 1993 and Bertolani *et al.* 2011a), *Milnesium tardigradum* (redescribed by Michalczyk *et al.* 2012), *Mesocrista spitsbergensis* (redescribed by Gąsiorek *et al.* 2016), *Hypsibius dujardini* (redescribed by Gąsiorek *et al.* 2018) or *Ramazzottius oberhaeuseri* (redescribed by Stec *et al.* 2018c). Thus, in our

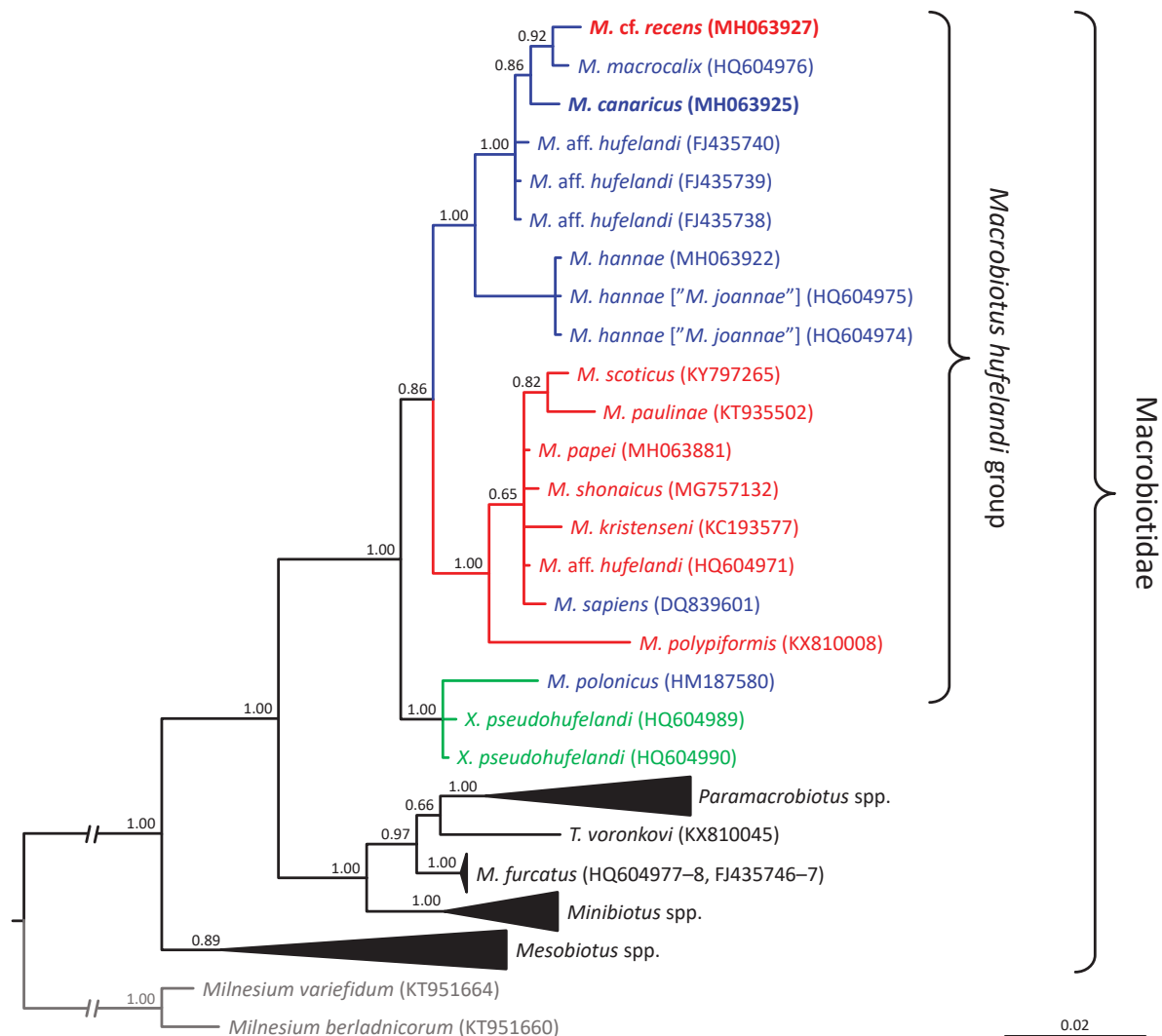
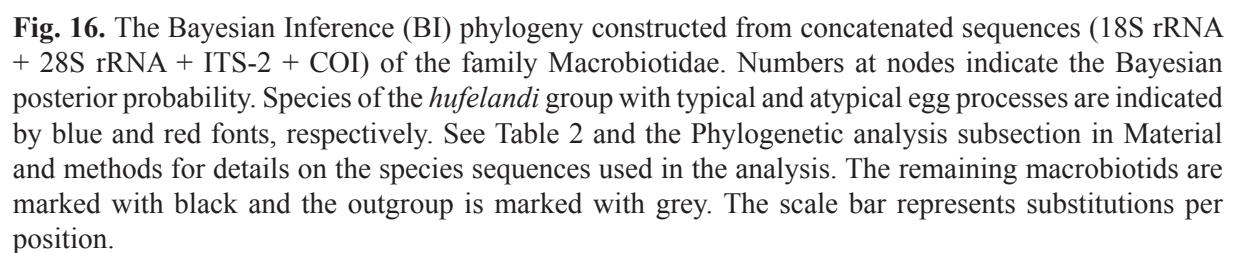


Fig. 15. The Bayesian Inference (BI) phylogeny constructed from 18S rRNA sequences of the family Macrobiotidae. Numbers at nodes indicate the Bayesian posterior probability. Species of the *hufelandi* group with typical and atypical egg processes are indicated by blue and red fonts, respectively. The *X. pseudohufelandi* + *M. polonicus* clade, which is in a sister relationship to all other species of the *hufelandi* group, is indicated by green branches. See Table 2 and the Phylogenetic analysis subsection in Material and methods for details on species sequences used in the analysis. The remaining macrobiotids are marked with black and the outgroup is marked with grey. Scale bar represents substitutions per position.

Recently, Stec *et al.* (2018a) showed two well supported COI lineages within the *M. hufelandi* group: one clade grouping species with *hufelandi*-type egg processes in the shape of inverted goblets, and the other with modified egg process (conical processes or processes with filaments growing out of terminal discs). Thus, *M. cf. recens*, with its conical egg processes, should be expected to cluster with species with modified egg processes. However, in contrast to this prediction, *M. cf. recens* is embedded within the clade with species exhibiting typical egg morphology in our COI analysis (Fig. 17). Moreover, the



same position of the species was recovered in the 18S rRNA phylogeny (Fig. 15) and in the analysis based on four concatenated markers (Fig. 16). Also, *M. sapiens*, with typical egg processes, should be present in the *hufelandi* clade, but in our analyses it clustered with species exhibiting mostly modified processes (Fig. 15). Importantly, however, the 18S rRNA sequence identified as *M. sapiens* (DQ839601; Schill & Steinbrück 2007) comes from Croatia, whereas the type locality of the species is in Sicily (Binda & Pilato 1984). Thus, given that there are no type/neotype sequences for *M. sapiens* and Schill & Steinbrück (2007) did not provide SEM photomicrographs of Croatian eggs, it is not possible to verify the identity of the DQ839601 sequence. In other words, the sequence may represent a similar species of the *hufelandi* group that may exhibit terminal disc filaments. Nevertheless, regardless of the phylogenetic position of *M. sapiens*, the affinity of *M. cf. recens* with species exhibiting typical egg processes shows that the morphological criterion proposed by Stec *et al.* (2018a) to distinguish the two clades is not

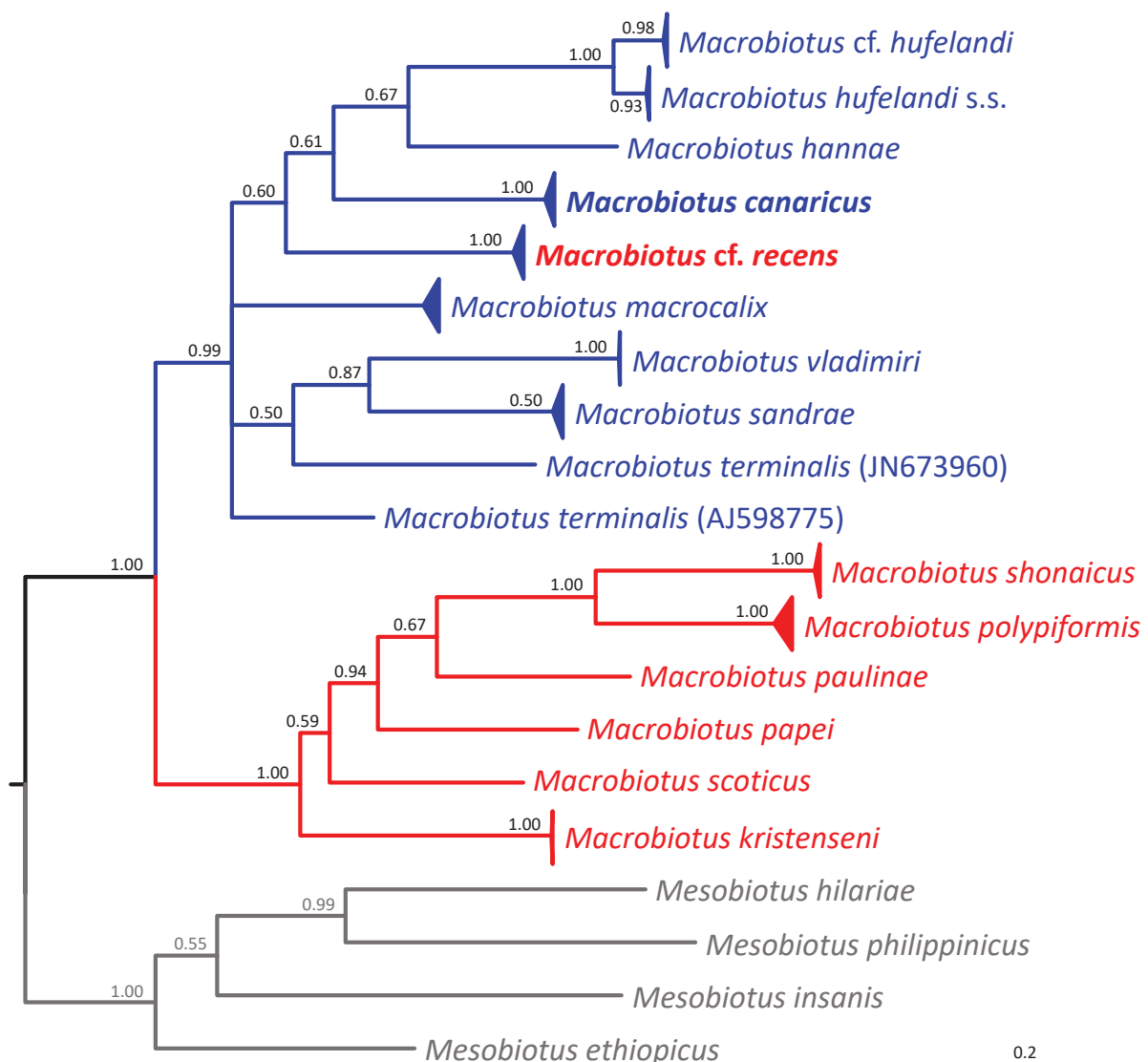


Fig. 17. The Bayesian Inference (BI) phylogeny constructed from COI sequences of the species of the *Macrobiotus hufelandi* group. Numbers at nodes indicate the Bayesian posterior probability. Species of the *hufelandi* group with typical and atypical egg processes are indicated by blue and red fonts, respectively. See Table 2 for details on the species sequences used in the analysis. The outgroup is marked with grey. The scale bar represents substitutions per position.

universal. Therefore, more species representing the two clades need to be sequenced to elucidate the taxonomic status of the two clades.

Another species whose phylogenetic position diverges from the predicted is *M. polonicus*. This species, with typical egg processes, should be embedded within the *hufelandi* clade. However, in the 18S rRNA phylogeny, not only does it not cluster within the *hufelandi* clade, but it clusters with *Xerobiotus pseudohufelandi* in a clade that is in a sister relationship to the entire *hufelandi* group. The *M. polonicus* + *X. pseudohufelandi* clade was also found to be a sister group to all other *hufelandi* group species by Bertolani *et al.* (2014). The genus *Xerobiotus* Bertolani & Biserov, 1996 shares a similar morphology of the buccal apparatus, egg shell and spermatozoa with species of the *M. hufelandi* group, but it differs from them by having strongly reduced claws. Given that the *M. polonicus* + *X. pseudohufelandi* clade contains only a single *hufelandi* group sequence, the clade could be a statistical artefact or possibly the 18S rRNA marker is too conservative to solve the relationships between the *hufelandi* group and the genus *Xerobiotus*. Therefore, it is crucial to analyse more populations of *M. polonicus*, other species of the *persimilis* subgroup (*sensu* Kaczmarek & Michalczyk 2017b) and species of the genus *Xerobiotus*, both in terms of the 18S rRNA marker and additional, more variable DNA fragments. If, however, more species of the *hufelandi* group turn out to cluster with the genus *Xerobiotus*, then the taxonomic status of the genus and of the *hufelandi* group should be reconsidered.

Although our multilocus phylogeny supports the presence of two distinct evolutionary lineages within the *hufelandi* group, the taxon sample size is still very low and more precise conclusions concerning the phylogeny of the group cannot currently be made. Thus, to provide more reliable conclusions, much more effort should be made to obtain multilocus molecular data linked to morphology for a larger number of species and populations. Particularly, it would be beneficial to increase the sample size, both in terms of markers as well as species, for the *M. polonicus* + *X. pseudohufelandi* clade to test whether the genus *Xerobiotus* and the *hufelandi* group are monophyletic. Another clade which would benefit from an increased sample size is the clade that comprises species with modified egg processes. Currently, the clade consists of species with two very different process morphotypes, i.e., species having processes with flexible filaments on the terminal discs and species having conical processes devoid of terminal discs. Thus, it might be possible that increased sampling could reveal the presence of two further clades that differ by the egg process morphology. Nevertheless, our findings support the previous results that egg morphology seems to be evolving faster than animal morphology, which underlines the usefulness of chorion ornamentation in the delineation of closely related species (Guidetti *et al.* 2013; Stec *et al.* 2016a, 2017a, 2018a).

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