UNRAVELLING THE TANGLE OF THE AZECID LAND SNAILS: A SURVEY ON THE SUPRASPECIFIC SYSTEMATICS BASED ON COMPARATIVE MORPHOLOGY AND MOLECULAR PHYLOGENY (GASTROPODA: EUPULMONATA: ORTHURETHRA)

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ABSTRACT: A selection of 17 azecid species, including Azeca goodalli, Gomeziella girottii, Hypnocarnica micaelae, four species of Cryptazeca and ten species currently assigned to Hypnophila, chosen among those best known, were investigated to elucidate their relationships based on morphological and molecular evidence. Thirty one characters, only 15 of which were parsimony informative, were scored from morphology. Parsimony analysis was performed with PAUP* 4.0 using Cochlicopa lubrica as outgroup. Sequences of COI and ITS2 deposited in GenBank were re-examined using Cochlicopa lubrica as outgroup. Phylogeny based on the morphological characters suggested that Hypnophila as formerly conceived was a polyphyletic taxon with four different lineages: Gomeziella girottii (1); Hypnophila boissii (2); the Dalmatian Hypnophila species except Hypnophila zirjensis (3); the western Mediterranean species plus Hypnophila zirjensis (4). Unfortunately molecular studies did not include Gomeziella girottii and species of Dalmatian Hypnophila and this biases full comparison between the two data sets. Moreover our re-analysis of the previous molecular data produced a slightly different phylogenetic hypothesis compared to the original one. Indeed, the only ML tree of COI sequences confirmed with weak bootstrap supported the phylogenetic hypothesis that Hypnocarnica micaelae was the sister group to all the other azecids. On the contrary, the ML trees of ITS2 and of concatenated COI+ITS2 sequences found that Azeca goodalli was the sister group to all the other azecids, Cryptazeca was paraphyletic and Hypnocarnica micaelae was the sister group of Cryptazeca monodonta. Comparing the results of the two data sets, it emerged that phylogenetic analysis based on morphological characters had good resolution but very low statistical support and that the position of Hypnophila boissii was the most variable. In morphological phylogeny this species had unresolved relationships with Gomeziella girottii and a clade including Azeca goodalli and all other species currently assigned to Hypnophila, whereas in the molecular phylogeny it was nested in the western Hypnophila. The latter result is followed here assigning the species now included in Hypnophila to two distinct genera: Gomphroa comprising the western species plus Hypnophila zirjensis; Hypnophila including the Dalmatian species except Hypnophila zirjensis.

KEY WORDS: Azecidae, morphological and molecular phylogeny, systematics, description of taxa
INTRODUCTION

Azecids are a group of small orthurethran land snails inhabiting the litter and topsoil of arid to humid open to forest environments. They occur in the western Palaeartic from the British Isles southward to the Iberian Peninsula and eastward to the western sector of the Balkan Peninsula in Europe and from Morocco to Algeria in North Africa (Holyoak & Holyoak 2012, Welter-Schultes 2012, Štampol et al. 2018).

The systematic history of this group of land snails is intricate, due to the fact that they have been regarded as belonging to the orthurethran cochlicopids or to the sigmurethran ferussaciids. A family-group taxon for this group (“Azicinae”) was first introduced by Watson (1920) as a subfamily of the Pupillidae to separate Azeca Fleming, 1828 from the usually considered closely allied Cochlicopa Férussac, 1821. Subsequently this family-group taxon was disregarded as a junior synonym of the Cochlicopidae (for the last examples see Schileyko 1998a, Barker 1999), only to be resurrected recently as a distinct family in the Cochlicopoidea (Bank et al. 2001, Falkner et al. 2002) or as a distinct subfamily in the Cochlicopidae (Bouchet & Rocroi 2005). Madeira et al. (2010) definitively demonstrated that the azecids were distinct from the cochlicopids and the ferussaciids: the azecids and all the other orthurethran groups examined belonged to the non-achatinoid clade, whereas the ferussaciids belonged to the sigmurethran achatinoid clade; within the orthurethrans, the azecids had unresolved relationships with the chondrinids and a large assemblage including all the other orthurethrans so far examined. After that the azecids were accepted as a distinct family within the orthurethran pulmonates (e.g. Holyoak & Holyoak 2012, Bank & Neubert 2017, Bouchet et al. 2017, Cianfanelli et al. 2018a, b, Štampol et al. 2018).

According to Bank et al. (2001) and Falkner et al. (2002), the azecids have only two genera, Azeca and Hypnophila Bourguignat, 1858. A third genus, Gomphroa Westerlund, 1902, is only recognised by Schileyko (1998a). A fourth genus, Cryptazeca Folini et Bérillon, 1877(a), initially assigned to the sigmurethran ferussaciids, was recently allocated to the azecids on the basis of morphological (orthurethrous kidney) and molecular evidence (Gómez & Angulo 1987, Madeira et al. 2010). A fifth genus, Gomezilla Cianfanelli, Bodon, Giusti et Manganelli, 2018(a), was established for Hypnophila girottii Eus, 1978, a species first described from the Plio-Pleistocene of Sardinia (western Mediterranean) and then discovered alive (Manganelli et al. 1995, Cianfanelli et al. 2018a). Finally a sixth genus, Hypnocarnica Cianfanelli et Bodon in Cianfanelli et al., 2018 was introduced for a new species (Hypnocarnica micaelaes Cianfanelli et Bodon in Cianfanelli et al., 2018) from the Carnic Prealps, Italy (Cianfanelli et al. 2018b).

Here we re-examine superspecific level relationships of the azecids. We conducted a phylogenetic study on the basis of morphological characters, considering a selection of the best known species, including the type species of all the genus-group taxa of the family. Then we compared the results with those of two molecular phylogenies, one from Cianfanelli et al. (2018b) and one based on a re-examination of the data of Cianfanelli et al. (2018b). Finally we made a concise survey of all the azecid genera, preceded by re-description of the family.

MATERIAL AND METHODS

TAXONOMIC SAMPLE

Our analysis considered a selection of azecid species including Azeca goodalli (Férussac, 1821), Gomeziella girottii, Hypnocarnica micaelaes, four species of Cryptazeca and ten species currently assigned to Hypnophila chosen among those best known.

OUTGROUP

Azecids have unresolved relationships with the chondrinids and a large assemblage including all the other orthurethran genera so far examined (Madeira et al. 2010). Under such uncertainty, we used Cochlicopa lubrica (Müller, 1774) (Figs 1–5), traditionally regarded as allied with the azecids, as outgroup.

MORPHOLOGICAL STUDY

For each species considered, detailed anatomical re-examination was performed on the available material. The bodies were isolated from the shells and dissected under the light microscope (Wild MSA) using very fine pointed watchmaker’s tweezers. Anatomical details were drawn using a Wild camera lucida.

When possible the material examined is listed as follows: locality, municipality and province names in parenthesis, UTM reference, collector(s), date, number of specimens and collection in which material is kept in parenthesis. Key to acronyms used in material examined: AMC – A. Margelli collection (Pisa, Italy); CGAH – D. T. Holyoak & G. A. Holyoak collection (Cabeço, Portugal); CNHM – Croatian
Figs 1–3. Shells of *Cochlicopa lubrica* (Müller, 1774): 1–2 – specimens from Mindelsee (09°02'E, 47°45'N) (district of Kostanz, Baden-Württemburg), G. Armbruster leg. 10.1992 (FGC 6439); 3 – a specimen from Mindelsee (09°01'E, 47°45'N) (district of Kostanz, Baden-Württemburg), G. Armbruster leg. 10.1992 (FGC 6440). Scale bar 2 mm

Figs 4–5. Genitalia of *Cochlicopa lubrica* (Müller, 1774): 4 – general view (gonad excluded); 5 – detail and internal structure of penial complex; a specimen from Tarvisio (province of Udine, Italy), F. Giusti leg. 4.9.1973. Scale bar 1 mm
Natural History Museum (Zagreb, Croatia); FGC – F. Giusti collection (Dipartimento di Scienze Fisiche, della Terra e dell’Ambiente, Università di Siena, Italy); MBC – M. Bodon collection (Genoa, Italy); MZB – Museu de Ciências Naturals (Barcelona, Spain); MZUF – Museo di Storia Naturale dell’Università di Firenze, Sezione Zoologica de “La Specola” (Florence, Italy); SCC – S. Cianfanelli collection (Firenze, Italy).

Key to acronyms used in figures: AG – albumen gland; BC – bursa copulatrix; BW – body wall; DBC – duct of bursa copulatrix; DDBC – diverticulum of duct of bursa copulatrix; E – epiphallus; FHD – first hermaphrodite duct; FO – free oviduct; GA – genital atrium; P – penis; PA – penial appendix; PD – penial diverticulum; POS – prostatic portion of ovispermic duct; PR – penial retractor muscle; PS – penial sheath; UOS – uterine portion of ovispermic duct; V – vagina; VM – vaginal muff. Other abbreviations in the text: D – shell maximum diameter; H – shell height; C, L, ML, M – radular teeth: central, lateral, latero-marginal and marginal, respectively.

In the shell description, apertural elements are defined: plicae or lamellae if they consist of elongated spiral structures situated in the palatum or parietum and columella respectively, teeth or denticles otherwise; the transversely elongated structure in the outermost parietum of many azecid species is named “angular tubercle” in line with Schileyko (1998a).

In the anatomical description the use of directional (proximal and distal; basal and apical) and descriptive terms (short vs. long, slender vs. wide, etc.) was applied as follows. In the case of the ducts of the main axis of the genitalia (e.g. free oviduct, vagina, vas deferens, penis, etc.), proximal, basal or initial denotes the part which is closer to the gonad and distal, apical, final or terminal the part which is closer to the gonopore. In the case of the blind structures radiating from the main axis of the genitalia (bursa copulatrix complex, penial/vaginal/atrial appendix, etc.), basal or initial denotes the part closer to, and apical, final or terminal, the part further from the main axis of the genitalia. The descriptive terms (e.g. short vs. long, slender vs. wide, etc.) refer to comparisons of the same tract of the genitalia in different taxa. Unfortunately, only extreme cases are easy to evaluate; intermediate cases are very difficult to evaluate objectively due to large variation between taxa. Application of these simple categories is straightforward. Apparently simple structures (penis or even bursa copulatrix, duct of bursa copulatrix, etc.) may be so varied in different taxa as to elude any attempt at objective comparison. Besides the variation between taxa, sometimes differences may be due to different sexual maturation of the reproductive organs, fixation, and in the case of drawings in the literature, different fidelity of illustration. Only examination of a considerable number of specimens by the same researcher can provide real insights, but this was not possible due to unavailability of material.

CHARACTERS

Thirty one characters were scored during examination of 17 azecid species belonging to the in-group, plus Cochlicopa lubrica used as out-group (Table 1). Eighteen characters were parsimony uninformative; only thirteen characters were parsimony informative.

PHYLOGENETIC ANALYSIS

Maximum parsimony analysis was performed with PAUP* 4.0b10 (Swofford 1998) with the following options: heuristic search mode with 1,000 random stepwise addition sequence replicates; tree bisection-reconnection branch swapping (TBR); MulTrees option “on”, saving all minimum trees found during branch swapping. All characters were considered unordered.

Once the initial heuristic search was completed, all the equally most-parsimonious trees were used for successive weighting to reduce the effect of potentially homoplastic characters (Farris 1969, Kores et al. 2000). Characters were assigned new weights using the Rescaled Consistency Index (RC) and a base weight of 100. Heuristic search was then performed on the reweighted matrix using branch swapping on all trees obtained in the previous analysis. After each search the characters were reweighted on the basis of the new trees and this procedure was repeated until a constant length was obtained. Multiple parsimonious trees were then summarised using a strict consensus method. To evaluate clad support, nonparametric bootstrap resampling with full heuristic search was used (1,000 bootstrap replicates, 10 random addition sequence replicates per bootstrap replicate, TBR swapping, MulTrees option “on”).

MOLECULAR ANALYSIS

Mitochondrial COI (cytochrome c oxidase subunit I) and 16S (16S ribosomal DNA) as well as nuclear ITS2 (internal transcribed spacer 2 of ribosomal DNA) partial gene sequences were obtained from GenBank. They were as follows: COI – MF545160 (Devaard 2017) and MG209139-MG209152 (Cianfanelli et al. 2018b); 16S – GU331954 (Dinapoli et al. 2011) and MG209153-MG209164 (Cianfanelli et al. 2018b); ITS2 – AY014019 (Wade et al. 2001), AY546470 (Armbruster et al. 2005), FJ791121-FJ791123 (Madeira et al. 2010), MG209165-MG209179 (Cianfanelli et al. 2018b).
Table 1. Description of characters used for morphological phylogenetic analyses

| No. | Name                          | Characters                                                                 | States                                                                 | Remarks                                                                                                                                                                                                                                                                                                                                 |
|-----|-------------------------------|----------------------------------------------------------------------------|----------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 1   | shell shape                   | 0 = elongate ovoid-fusiform (*Cochlicopa* and *Azeca* like)                 | 1 = elongate ovoid-cylindrical (*Hypnophila* like)                    | It is very difficult to propose an objective coding of shell shape variation due to significant intraspecies variability.                                                                                                                                                                                                                 |
|     |                               | 2 = elongate cylindrical-fusiform (*Gomphrea* like)                        | 3 = very elongate, cylindrical-conical or cylindrical-fusiform (*Gomeziella* like) |                                                                                                                                                                                                                                                                                                                                                                                                  |
|     |                               | 4 = very elongate, cylindrical-conical or cylindrical-fusiform (*Hypnocarnica* like) |                                                                      |                                                                                                                                                                                                                                                                                                                                                                                                  |
| 2   | whorls and sutures            | 0 = whorls slightly round and sutures deep                                 | 1 = whorls almost flat and sutures shallow                           | State 0 is unique to *Cochlicopa lubrica* (Müller, 1774).                                                                                                                                                                                                                                                                                                                                      |
| 3   | angular tubercle              | 0 = absent                                                                 | 1 = present                                                           | A transversely elongate tubercle is present in the outermost parietum of *Azeca goodalli* (Férussac, 1821) and most *Hypnophila* species ("triangular tooth near the upper angle" according to PILSBRY 1908; "angular tubercle" according to SCHELEYKO 1998a). State 1 is unique to *A. goodalli*.                                                                                      |
| 4   | supraparietal denticle/knob/tubercle | 0 = absent                                                                 | 1 = present                                                          | State 1 is unique to *A. goodalli*. Sometimes it is joined anteriorly/abapically to an infraparietal lamella via a transverse bridge (so that the two lamellae form a horseshoe-/U-shaped structure).                                                                                                                                                         |
| 5   | parietal lamella              | 0 = absent                                                                 | 1 = present                                                           | An infraparietal lamella is absent or present in *A. goodalli* and absent in all the other species. Sometimes it is joined anteriorly/abapically to a parietal lamella via a transverse bridge (so that the two lamellae form a horseshoe-/U-shaped structure). State 1 is unique to *A. goodalli*.                                                                 |
| 6   | infraparietal lamella         | 0 = absent                                                                 | 1 = present                                                           | A transversely elongate tubercle is sometimes present in the innermost parietum of *Gomeziella girottii* (Esu, 1978).                                                                                                                                                                                                                  |
| 7   | parietal tubercle             | 0 = absent                                                                 | 1 = present                                                           | State 1 is unique to *A. goodalli*.                                                                                                                                                                                                                                                                                               |
| 8   | columellar lamella            | 0 = absent                                                                 | 1 = present                                                           | The coding of the character was very difficult. A subcolumellar lamella is absent in *C. lubrica* and very evident in *A. goodalli* but these two states are the extremes of a continuum and many species show wide internal variability.                                                                                                                                         |
| 9   | subcolumellar lamella         | 0 = absent or only slightly evident                                       | 1 = evident                                                           | A lower palatal denticle/knob/tubercle is absent or present in *A. goodalli* and absent in all the other species.                                                                                                                                                                                                                     |
|     |                               | 2 = clearly evident                                                       |                                                                       | State 1 is present in *A. goodalli*. In *G. girottii*, at the transition point between the thin and thickened outer peristomal margin, there is sometimes a very subtle knob (in the former species it is clearly evident, whereas in the latter it is faint).                                                                       |
| 10  | lower palatal knob (tooth-like plica) | 0 = absent                                                                 | 1 = present                                                           | The coding of the character was very difficult. The outer peristomal margin in lateral view is almost straight in *Hypnophila* species and very sinuous in *A. goodalli* but between these two states many situations are subjective and some species show wide internal variability.                                                                                                                   |
| 11  | upper palatal knob (tooth-like plica) | 0 = absent                                                                 | 1 = present                                                           |                                                                                               |
| 12  | upper third of the outer peristomal margin | 0 = thickened as elsewhere                                                  | 1 = thinner than elsewhere                                            |                                                                                                                                                                                                                                                                                                                                                                                                |
| 13  | outer peristomal margin in lateral view | 0 = almost straight (only slightly sinuous)                                | 1 = sinuous to very sinuous and inverted S-like                        |                                                                                                                                                                                                                                                                                                                                                                                                |
| No. | Name | States | Remarks |
|-----|------|--------|---------|
| 14. | rows of pits on protoconch | 0 = absent | State 1 is unique to *A. goodalli*. |
| 15. | eye spots | 0 = absent | State 0 is unique to *C. lubrica*. |
| 16. | latero-marginal teeth | 0 = absent | State 0 is unique to *C. lubrica*. |
| 17. | diverticulum of bursa copulatrix duct | 0 = absent | The calibre of the vas deferens is seldom uniform; however only in *Cryptazeca vasconica* (Kobelt, 1894) and *G. girottii* is there a clearly evident medial thickening. |
| 18. | vaginal muff | 0 = absent | State 0 is unique to *C. lubrica*. |
| 19. | vas deferens | 0 = without evident medial thickening | State 0 is unique to *C. lubrica*. |
| 20. | epiphallus/vas deferens | 0 = enters penis basally | State 1 is unique to *Cryptazeca monodonta* (De Folin et Bérillon, 1877a). |
| 21. | epiphallus | 0 = present | The only slightly enlarged distal portion of vas deferens in *Azeca* and some *Cryptazeca* species has sometimes been regarded as an epiphallus (e.g. HESSE 1922, GOMEZ 1988, 1991). |
| 22. | penis | 0 = medium-sized or large, long and cylindrical | State 2 is unique to *Hypnophila boissii* (Dupuy, 1851) and state 3 to *C. monodonta*. |
| 23. | muscular sheath on proximal penis | 0 = absent | State 1 is unique to *A. goodalli*. |
| 24. | penial appendix | 0 = absent | State 0 is unique to *C. lubrica*. |
| 25. | penial diverticulum | 0 = absent | A penial diverticulum is absent or present in *A. goodalli* and absent in all the other species. |
| 26. | penial retractor | 0 = unbranched | The penial retractor is unbranched or branched in *A. goodalli* and unbranched in all the other species. |
| 27. | position of penial retractor | 0 = lateral | State 0 is unique to *C. lubrica* and *G. girottii*. |
| 28. | internal structures at epiphallus/vas deferens opening into penis | 0 = conical papilla | |
| 29. | internal penial structures (longitudinal pleats) | 0 = absent | We assumed homology of the tongue-like structures/bulges/ring-shaped sarcobelum inside the proximal penis of *Cryptazeca* species and *Hypnocarnica micaela* Cianfanelli et Bodon in Cianfanelli et al., 2018(b) based on their shape and location. |
| 30. | internal penial structures (thorned papillae) | 0 = absent | |
| 31. | internal penial structures (tongue-like structure inside proximal penis) | 0 = absent | |

Table 1. continued
The alignments were performed using the CLUSTAL W programme (Thompson et al. 1994) implemented in BIOEDIT, version 7.0.5 (Hall 1999). The COI sequences were aligned according to the translated amino acid sequences. The ends of all sequences were trimmed. The lengths of the sequences after cutting were 588 bp for COI, 358 positions for 16S and 748 positions for ITS2. Concatenated COI+16S sequences were of 1,396 positions long (615 COI + 781 ITS2).

Maximum Likelihood (ML) analyses were performed with MEGA 7 (Kumar et al. 2016). For each alignment file the best nucleotide substitution model was specified according to the Bayesian Information Criterion (BIC) (Tamura 1992, Kumar et al. 2016): T92+G+I for COI, ITS2 or COI+ITS2 concatenated sequences and T92+G for 16S. In parallel, pairwise distances based on COI sequences were calculated using the Kimura two-parameter model (K2P) (Kimura 1980).

RESULTS AND DISCUSSION

AZECID PHYLOGENY BASED ON MORPHOLOGICAL CHARACTERS

Phylogenetic analysis (Fig. 6) had good resolution (only 9 trees with length L = 2824; consistency index CI = 0.95; retention index RI = 0.95; rescaled consistency index RC = 0.91) but statistical support was very low (only seven clades were supported, five of them very weakly).

Phylogenetic analysis (Tables 2 and 3) found that most of the species belong to a main clade (MAC – main azecid clade) including Azeca goodalli and all species currently assigned to Hypnophila except Hypnophila boissii (Dupuy, 1851). The nine trees differ in the relationships between MAC, Gomeziella girottii and H. boissii (Table 3) and in the relationships between Cryptazeca species. In the consensus tree MAC has unresolved relationships with G. girottii and H. boissii. In turn MAC includes two subclades: one with A. goodalli and the Dalmatian Hypnophila species except Hypnophila zirjensis Štamol, Manganelli, Barbaro et Giusti, 2018, the other comprising the remaining western Mediterranean Hypnophila species plus H. zirjensis. Finally, the group comprising MAC, G. girottii and H. boissii has sister group relationships with Cryptazeca species; in turn this larger group has sister group relationships with Hypnocarnica micaelae.

AZECID PHYLOGENY BASED ON MOLECULAR SEQUENCES

Cianfanelli et al. (2018b) published a Bayesian Inference tree of concatenated dataset of three genes (COI, 16S, ITS2) for 16 azecids representative of 12 species (Azeca goodalli, Cryptazeca monodonta (De Folin et Bérillon, 1877(a)), C. spelaea Gómez, 1990(c), Hypnophila bisacchii Giusti, 1970, H. boissii, H. dohrni Paulucci, 1882, H. etrusca Paulucci, 1886, H. malagana Gittenberger et Menkhorst in Gittenberger, 1983, H. remyi Boettger, 1949, Hypnophila sp. A, Hypnophila sp. B, Hypnocarnica micaelae). Their tree of concatenated sequences COI+16S+5.8S+ITS2+28S
Table 2. Character-taxon matrix used for phylogenetic analyses

| Taxa/Characters | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 | 24 | 25 | 26 | 27 | 28 | 29 | 30 | 31 |
|----------------|---|---|---|---|---|---|---|---|---|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|
| Cochlicopa ulbricata | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Azeca goodalli | 0 | 1 | 1 | 1 | 1 | 0,1 | 0 | 1 | 1 | 0,1 | 1 | 1 | 1 | 1 | 1 | 1 | 0 | 0 | 1 | 1 | 1 | 0 | 0 | 1 | 1 | 1 | 0,1 | 1 | 1 | 1 | 3 | 1 | 0 |
| Cryptazeca elongata | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0,1 | 0 | 0 | 1 | 1 | 1 | ? | ? | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 1 | 3 | 0 | 1 | 2 |
| Cryptazeca monononta | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 1 | 0 | ? | 1 | 1 | 0 | 1 | 1 | 3 | 0 | 1 | 1 | 2 |
| Cryptazeca speleae | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0,1 | 0 | 0 | 1 | 1 | 1 | ? | 0 | 1 | 1 | 0 | 1 | 0 | 0 | 1 | 0 | 1 | 0 | 1 | 3 | 0 | 1 | 2 |
| Cryptazeca vasconia | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0,1 | 0 | 0 | 1 | 1 | 1 | ? | ? | 1 | 1 | 1 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 1 | 3 | 0 | 1 |
| Gomeziella gigoti | 2 | 1 | 0 | 0 | 0 | 0,1 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 1 | 0 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 3 | 1 | 0 |
| Hypnophila pusapiformis | 3 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 1 | 1 | 0 |
| Hypnophila zacynthia | 3 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | ? | ? | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 1 | 1 | 0 |
| Hypnophila bisachi | 0 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila boissii | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila dohini | 1 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila emiliana | 1 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila etrusca | 1 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 1 | 0 | 1 | 1 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila incerta | 1 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila remy | 1 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 1 | 2 | 1 | 0 |
| Hypnophila zirjensis | 1 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | ? | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 1 | 2 | 1 | 0 |
| Hypnocarnica miaeae | 4 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0,1 | 0 | 0 | 1 | 1 | 1 | 0 | 1 | 1 | 0 | 0 | 1 | 0 | 0 | 1 | 3 | 0 | 0 | 1 |

Polymorphic states (6, 10, 25, 26 in Azeca goodalli; 7 in Gomeziella gigoti; 9 in Cryptazeca elongata, C. speleae and C. vasconia) were treated as unknown (?) in the analysis.

Table 3. List of main lineages and their auto/synapomorphies

| Clades | trees 1, 2, 3 | trees 4, 5, 6 | trees 7, 8, 9 |
|--------|---------------|---------------|---------------|
| Azecids | (MAC, BOI, GIR) | (MAC, BOI, GIR) | (MAC, BOI, GIR) |
| Hypnocarnica miaeae | 1(4), 15(1), 31(1) | 1(4), 15(1), 31(1) | 1(4), 15(1), 31(1) |
| Cryptazeca + Gomeziella gigoti + Hypnophila boissii + MAC | 12(1), 28(3) | 12(1), 28(3) | 12(1), 28(3) |
| Cryptazeca | 30(1), 31(2) | 13(1), 30(1), 31(2) | 13(1), 30(1), 31(2) |
| Gomeziella gigoti + Hypnophila boissii + MAC | 29(1) | 29(1) | 29(1) |
| Hypnophila boissii + MAC | 13(0) | 13(0) | 13(0) |
| Gomeziella gigoti + Hypnophila boissii | 1(1) | 1(1) | 1(1) |
| Gomeziella gigoti | 1(3), 15(1), 19(1), 27(0) | 1(3), 15(1), 19(1), 27(0) | 1(3), 15(1), 19(1), 27(0) |
| Hypnophila boissii | 1(1), 22(2) | 22(2) | 1(1), 22(2) |
| MAC | 3(1) | 3(1) | 3(1) |
| Azeca + Dalmatian Hypnophila except Hypnophila zirjensis | 14(0) | 14(0) | 14(0) |
| Azeca | 4(1), 5(1), 8(1), 11(1), 12(1), 16(1), 23(1) | 4(1), 5(1), 8(1), 11(1), 12(1), 16(1), 23(1) | 4(1), 5(1), 8(1), 11(1), 12(1), 16(1), 23(1) |
| Dalmatian Hypnophila except Hypnophila zirjensis | 1(2), 28(1) | 1(2), 28(1) | 1(2), 28(1) |
| Western Hypnophila plus Hypnophila zirjensis | 28(2) | 28(2) | 28(2) |

BOI – Hypnophila boissii, GIR – Gomeziella gigoti, MAC – main azecid clade including Azeca and species currently assigned to Hypnophila except H. boissii.
(Cianfanelli et al. 2018b: fig. 7) shows phylogenetic relationship of 16 taxa but the list of sequences deposited in GenBank (Cianfanelli et al. 2018b: tab. 1) contains full set of three genes for only 11 taxa. GenBank resources on azecids are very scanty. Apart from the sequences deposited by Cianfanelli et al. (2018b) there are only a few sequences of ITS2 deposited by Armarbuster et al. (2005) and Madeira et al. (2010). Therefore, we re-examined separately the sequences for COI, 16S and ITS2 deposited in GenBank by Cianfanelli et al. (2018b) and some ITS2 deposited by Madeira et al. (2010) using specified sequences of Cochlicopa lubrica as outgroup. Our separate analyses for each gene confirmed that all 12 taxa form one clade in ML trees, as shown by our analysis of morphological characters, whereas the separation of subclades differs for particular genes.

We do not show the Maximum Likelihood (ML) tree of 16S sequences because Cianfanelli et al. (2018b) did not deposit four sequences of species analysed (i.e. Cryptazeca spelaea 1, C. spelaea 2, Hypnophila malagana 1, H. malagana 2) in GenBank.

Molecular phylogenetic analysis based on COI sequences was performed with only 14 COI sequences (Fig. 7) because there were no GenBank deposits for A. goodalli 1 and C. spelaea 2 (Cianfanelli et al. 2018b: tab. 1). Azeca goodalli seems to be very different from all other azecids and forms a separate subclade. K2P genetic distances between COI sequences indicate that A. goodalli differs by 29.1–35.6% from the species representing three other azecid genera (Table 4). Separateness of Hypnocarnica micaelae is also visible (as its K2P values distance it by 21.2–35.6% from other species). All other sequences cluster together in three further subclades, two for species included in Hypnophila and one for Cryptazeca. K2P distances between COI sequences of species of these genera, i.e. Cryptazeca vs. Hypnophila, Cryptazeca vs. Hypnocarnica and Hypnophila vs. Hypnocarnica, are smaller (16.3–26.4%, Table 4), however they support attribution of particular species to different genera. It is noteworthy that the K2P distance between two different specimens of Hypnophila malagana originating from two different Spanish populations is similarly large (16.2%). There is also a very large K2P distance (22.5%) between MG209144 Hypnophila boissii and MG209145 Hypnophila sp. A (in GenBank named Hypnophila boissii), i.e. this large distance differentiates sequences of French (MG209144) and Spanish

Table 4. K2P genetic distances between analysed COI sequences

|                  | K2P distance (%) |
|------------------|------------------|
| Within Azeca     | 0.0              |
| Within Cryptazeca| 10.4–17.4        |
| Within Gomphroa  | 8.5–22.4         |
| Within Hypnocarnica| 0.0            |
| Azeca vs Cryptazeca| 31.2–32.2       |
| Azeca vs Gomphroa| 29.1–32.8        |
| Azeca vs Hypnocarnica| 35.6          |
| Cryptazeca vs Gomphroa| 16.3–24.9     |
| Cryptazeca vs Hypnocarnica| 21.2–24.9   |
| Gomphroa vs Hypnocarnica| 22.6–26.4    |

We do not show the Maximum Likelihood (ML) tree of 16S sequences because Cianfanelli et al. (2018b) did not deposit four sequences of species analysed (i.e. Cryptazeca spelaea 1, C. spelaea 2, Hypnophila malagana 1, H. malagana 2) in GenBank.

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| Azeca vs Gomphroa| 29.1–32.8        |
| Azeca vs Hypnocarnica| 35.6          |
| Cryptazeca vs Gomphroa| 16.3–24.9     |
| Cryptazeca vs Hypnocarnica| 21.2–24.9   |
| Gomphroa vs Hypnocarnica| 22.6–26.4    |

We do not show the Maximum Likelihood (ML) tree of 16S sequences because Cianfanelli et al. (2018b) did not deposit four sequences of species analysed (i.e. Cryptazeca spelaea 1, C. spelaea 2, Hypnophila malagana 1, H. malagana 2) in GenBank.

Molecular phylogenetic analysis based on COI sequences was performed with only 14 COI sequences (Fig. 7) because there were no GenBank deposits for A. goodalli 1 and C. spelaea 2 (Cianfanelli et al. 2018b: tab. 1). Azeca goodalli seems to be very different from all other azecids and forms a separate subclade. K2P genetic distances between COI sequences indicate that A. goodalli differs by 29.1–35.6% from the species representing three other azecid genera (Table 4). Separateness of Hypnocarnica micaelae is also visible (as its K2P values distance it by 21.2–35.6% from other species). All other sequences cluster together in three further subclades, two for species included in Hypnophila and one for Cryptazeca. K2P distances between COI sequences of species of these genera, i.e. Cryptazeca vs. Hypnophila, Cryptazeca vs. Hypnocarnica and Hypnophila vs. Hypnocarnica, are smaller (16.3–26.4%, Table 4), however they support attribution of particular species to different genera. It is noteworthy that the K2P distance between two different specimens of Hypnophila malagana originating from two different Spanish populations is similarly large (16.2%). There is also a very large K2P distance (22.5%) between MG209144 Hypnophila boissii and MG209145 Hypnophila sp. A (in GenBank named Hypnophila boissii), i.e. this large distance differentiates sequences of French (MG209144) and Spanish
(MG209145) populations of H. boissii. Our results therefore support the suggestion of Cianfanelli et al. (2018b) that French populations of H. boissii belong to a different species (probably Hypnophila etrusca or a species very close to it). The same is suggested by our analysis of populations of H. malagana. Of course, these suggestions must be verified in further anatomical and molecular studies involving a much larger number of individuals of all the populations involved.

The ML tree of ITS2 (Fig. 8) was based on 15 sequences deposited in GenBank by Cianfanelli et al. (2018b, the sequence for Hypnophila malagana 1 is lacking in their table 1). Using four other sequences (AY546470 and FJ791121-FJ791123, obtained from GenBank) we found that a clade with Azeca goodalli is separate from the other azecid species in question. Moreover, the sequences from species included in populations of Hypnophila from the western Mediterranean clustered separately from those included in Cryptazeca and Hypnocarnica. The lack of materials prevented study of the molecular relationships between Gomeziella and Hypnophila from the eastern Mediterranean and other azecid genera.

The ML tree of COI and ITS2 (Fig. 9) was based on these sequences deposited in GenBank by Cianfanelli et al. (2018b) which allowed to create concatenated COI+ITS2 sequences. The result was similar. Azeca was on a very different branch, Hypnophila from the western Mediterranean populations clustered separately from those included in Cryptazeca and Hypnocarnica.

Two final remarks: first, as we stated previously (Pieńkowska et al. 2019), molecular data alone cannot be used in taxonomic research. Molecular analysis must be supported by morphological features (of shells and/or genital anatomy) before any decision is made about taxonomy or nomenclature. Secondly, mitochondrial COI and 16S as well as nuclear ITS2 gene fragments belong to very variable parts of the genome, and are therefore suitable for studying molecular variation at population or species level. For studies at order or family level, less variable genes are more useful, which is why Madeira et al. (2010) excluded ITS2 sequences when studying the position of azecids in the Stylommatophora. They comment as follows: “Sequence variation in the ITS2 region was extremely high, and most of its sites could not be

Fig. 8. Maximum Likelihood (ML) tree of ITS2 sequences of Azecidae, based on sequences obtained from GenBank: AY546470 deposited by Armbruster et al. (2005), FJ791121-FJ791123 by Madeira et al. (2010) and MG209165-MG209178 by Cianfanelli et al. (2018b). Numbers next to the branches indicate bootstrap support above 50% calculated for 1,000 replicates (Felsenstein 1985). The tree was rooted with Cochlicopa lubrica sequence AY014019 deposited in GenBank (Wade et al. 2001).
Supraspecific systematics of Azecidae

AZECID SYSTEMATICS: THE STATE OF THE ART

Phylogeny based on morphological characters (Fig. 6) suggests that Hypnophila as formerly conceived is a polyphyletic taxon that includes four different lineages: Gomeziella girottii (1), Hypnophila boissii (2), the Dalmatian Hypnophila species except H. zirjensis (3); the western Mediterranean species plus H. zirjensis (4). Unfortunately molecular studies did not include any species of Dalmatian Hypnophila and Gomeziella girottii and this biases full comparisons between the two data sets. Moreover, our re-analysis of the molecular data by Cianfanelli et al. (2018b) produced a somewhat different phylogenetic hypothesis with respect to the original one. Indeed, only the ML tree of COI sequences (Fig. 7) confirmed (with weak bootstrap support) the phylogenetic hypothesis of Cianfanelli et al. (2018b) that Hypnocarnica micaelae was the sister group to all the other azecids. On the contrary, the ML tree of ITS2 sequences (Fig. 8) as well as the ML tree of concatenated COI+ITS2 sequences (Fig. 9) showed that Azeca goodalli was the sister group of all the other azecids, Cryptazeca was paraphyletic and Hypnocarnica micaelae was the sister group of Cryptazeca monodonta. In all the phylogenies, western Hypnophila species usually belonged to the same clade, but their internal relationships were variable.

Comparing the results of the two data sets, it emerges that phylogenetic analysis based on morphological characters has good resolution but very low statistical support and that the position of Hypnophila boissii is the most variable. This species has unresolved relationships with the main azecid clade and Gomeziella girottii in the morphological phylogeny, whereas it is nested in the western Hypnophila in the molecular phylogeny.

We think that our molecular results allow a new interpretation of morphological results, therefore we assigned the species now included in Hypnophila to two distinct genera: Gomphroa including the western species plus H. zirjensis; Hypnophila including the Dalmatian species except H. zirjensis.
SYSTEMATIC REVISION

This section includes a concise survey of all the azecid genera, preceded by a re-description of the family. The genera are listed alphabetically. For each taxon a short diagnosis, a detailed description of the type species and some remarks are given. At the end we also give a re-description of Hypnocarnica micaelae followed by some remarks on its systematics.

Family: Azecidae Watson, 1920
Type genus: Azeca Fleming, 1828

Diagnosis
Orthurethran land snails, according to phylogeny based on morphology, characterised by dextral, small, imperforate, elongate shell with whorls almost flat and sutures shallow (2(1)) and rows of pits on protoconch (14(1)); distal genitalia without diverticulum of bursa copulatrix duct (17(1)), epiphallus (21(1)) or penial appendix (24(1)) and with vaginal muff (18(1)) and basal/sub-basal insertion of penial retractor (27(1)). Some taxa lack rows of pits on the protoconch and have additional structures, which replace the true penial papilla, bordering where the vas deferens opens into the penis.

Description
Body: snail-like.
Foot: holopod, undivided.
Jaw: aulacognathe.
Radula: central tooth small, tricuspid; lateral teeth large, bicuspid; latero-marginal teeth absent (Azeca) or present (tricuspid, because ectocone split into two); marginal teeth pluricuspid.
Shell: dextral, small (shell height less than 10 mm), imperforate, elongate to very elongate, ovoid-fusiform, ovoid-cylindrical, cylindrical-fusiform, cylindrical-conical, pale-brown, reddish-brown, yellowish or colourless, glossy and transparent when fresh, with 5 to 10 slightly convex to flat whorls, separated by superficial sutures; aperture orthocline to slightly prosocone, oblique pyriform to ovate pyriform, pointed to rounded at base, with apertural armature very complex to simplified or absent (up to 7–8 pieces: supraparietal, parietal, infraparietal, columellar and subcolumellar lamellae; one or two lower palatal and one upper palatal tooth-like plicae); peristome not reflected, thin in its outer upper third, slightly thickened or thickened elsewhere, sometimes with callous rim on columella and parietum ending in transversely elongate angular tubercle separated by deep notch from upper angle/vertex of aperture and with straight or sinuous (more or less inverted S-like) outer margin; when parietal lamella is present, angular tubercle and upper palatal knob delimit more or less evident sinulus (variably deep furrow may occur at upper angle of aperture between angular tubercle and outer peristome); protoconch smooth or with faint radial crests or close spiral rows of small pits; teleconch smooth, with very thin spiral grooves and sometimes weak collabral growth lines.

Proximal genitalia (those including parts close to the gonad): the first hermaphroditic duct, talon (seminal receptacle – fertilisation pouch complex), albumen gland and second hermaphroditic duct (or ovispermduct) do not show significant variation between the various taxa, whereas distal genitalia have very complex and diverse structure.

Female distal genitalia include: free oviduct, bursa copulatrix and its duct and vagina. Free oviduct very short to long and variably wide. Bursa copulatrix sac-like, bean-like, oval or pyriform with long and slender to short and wide duct, initially not flared to very flared. Vagina short to long, variably wide, completely or partially covered by faint to thick, yellowish to brownish muff of “pigmentary cells” (according to GóMez 1991); internal surface smooth or with protuberance covered in small papillae with apical thorns (vaginal part of auxiliary copulatory organ, according to GóMez 1991).

Male distal genitalia include vas deferens and penis. Vas deferens rather long (shorter than FO + V + P) to very long (longer than FO + V + P), more or less uniform in diameter, entering penis basally, sub-basally, laterally or medially; in Azeca and Cryptazeca, slightly thickened in its proximal and subdistal tract; in Gomeziella thickened in its medial section; opening of vas deferens into penis simple (penial papilla absent) or bordered by initial part of one penial pleat or by a papilla consisting of bunch of small, conical appendices fused at base or by a true penial papilla. Penis very short to long, ovoid to pear-shaped or sub-cylindrical or uniformly cylindrical, undivided or divided into distinct proximal and distal parts by slight constriction (Azeca and Hypnocarnica). In Azeca proximal part of penis with or without penial diverticulum and with penial sheath. Penial diverticulum (Azeca), when present, very small to small, variable in shape and size (sac-like, elongate straight or hook-like). Penial sheath (Azeca) muff-like, enveloping final section of vas deferens and proximal penis, proximally usually continuous with penial retractor and distally ending level with slight constriction; sometimes part of proximal penis and its diverticulum (when present) protrude out of it. Penial retractor arising from diaphragm, inserted basally and sometimes enveloping final section of vas deferens (in Azeca continuous with penial sheath and sometimes with thin branch inserted on apex of penial diverticulum) or inserted basally/sub-basally close to or far from vas...
deferens or inserted laterally. Internal surface of penis with very small or large longitudinal pleats and/or spongy cords and accessory structures (thickened parts, knobs, transverse or longitudinal crests) or almost completely covered in small papillae with apical thorns.

Remarks

Within the geographical range of the family, only the orthurethran cochlicopids have a shell similar in shape and size to that of the azecids excluding Hypnocarnica micaelae. The latter has a shell very similar in shape, size and variation to that of the sigmurethran ferussacid Cecilioides Férussac, 1814, with very slender shells like those of Cecilioides acicula (Müller, 1774) and less slender shells like those of C. pettiana (Benoit, 1862) / C. tumulorum (Bourguignat, 1856) / C. janii (De Betta et Martinati, 1855). Again, all the azecids apart from Hypnocarnica micaelae always have the upper third of the outer peristome thinner than elsewhere. This feature is also detectable in species without a thickened peristome, such as Gomphroa boissii. Moreover, many azecids have at least one or two apertural folds/knobs inside the aperture. Azeca, Hypnophila and Gomphroa (excluding Gomphroa boissii) also have an angular tubercle. This transversely elongate apertural tubercle is the outermost element of the callous rim which borders the parietum and columella; it is separated from upper angle/vertex of aperture by a deep notch and is not related to the angular lamella (which joins the upper vertex/angle of the outer peristome).

Compared to other orthurethran groups, the azecids have simplified distal genitalia because they lack the diverticulum of the bursa copulatrix duct, the penial appendix and consequently the branch of the penial retractor attached to the penial appendix. The diverticulum of the bursa copulatrix duct and the penial appendix are absent in some unrelated groups of orthurethran snails (families Chondrinidae, Pyramidulidae, Pleurodiscidae, Vertiginidae and so on; Schileyko 1998a, b) and their absence may be regarded as a derived state (Baker 1935). The presence of the epiphallus is reported in Azeca and Cryptazeca by Hesse (1922) and Gómez (1988, 1991). However, the difference in calibre between the proximal and distal sections of the duct connecting the prostastic ovispermiduct to the penis is really minimal.

The structure of the ovispermiduct is also controversial. Gómez & Angulo (1990) and Gómez (1991) state that the female ovispermiduct of Azeca has an “oviductal caecum” in its proximal region and that the ovispermiduct of Azeca and Cryptazeca has a “blind-ending duct” in its proximal region, considered to be homologous with the allospermiduct of the other stylommatophoran species (Gómez & Angulo 1990: 109, 110, Gómez 1991: 99). The blind-ending duct was assumed by Gómez & Angulo (1990) to correspond to the diverticulum described by Boycott (1919). Our light microscope (Wild M5A) study of azecid anatomy failed to find the blind-ending duct (likewise F. Wiegmanns and P. Hesse; see Hesse (1922)) and the oviduct caecum in any of the taxa examined.

The more special anatomical features of azecids are the submedially thickened vas deferens of Gomeziella giriotii, the penial sheath of Azeca goodalli and the stimulator of Cryptazeca species. A submedially thickened vas deferens is known in a few other unrelated taxa (e.g. orculids: Alvariella Hausdorf, 1996 and Sphyradium Charpentier, 1837; see Hausdorf 1996). Penial sheaths are not identified in other orthurethrans, but this must be considered with caution, because they may have escaped detection during dissection. Finally, the stimulator of Cryptazeca, consisting of a vaginal and a penial bulge completely covered in small papillae with apically hooked thorns in the distal genitalia is unique, since similar structures are absent in other orthurethrans (Gómez 1991).

Genus: Azeca Fleming, 1828

Figs 10–28

Type species: Azeca tridens Fleming, 1828, by monotypy. Azeca tridens is a junior synonym of Helix goodalli Féruassac, 1821. According to ICZN (1999, Art. 11.10), the type species is Azeca tridens Fleming, 1828 and not Turbo tridens Pulteney, 1799 as stated by Opinion 335 (ICZN, 1953). The type species was indeed established by deliberate misidentification (according to Welter-Schultes 2012: Helix tridens Müller, 1774 as misidentified by Pulteney (1799)).

Material examined

Azeca goodalli (Férussac, 1821) – France, Nouvelle-Aquitaine: Pic de Gillers (department of Pyrénées-Atlantiques), 30TXN8463, B. Gómez leg. 30.10.1987 (4 spirit specimens, 3 dissected, FGC 41647); Sainte-Engrâce (department of Pyrénées-Atlantiques), 30TXN7762, B. Gómez leg. 28.7.1982 (2 spirit specimens, 1 dissected, FGC 41649); Bretagne: Louannec (department of Côtes-D’Armor), 30UVV60, M. Calcagnino & S. Cianfanelli leg. 30.6.1993 (4 spirit specimens, 3 dissected, SCC 18731); Occitania: quarry east of Lourdes (department of Hautes-Pyrénées), 30TYN4275, E. Bodon & M. Bodon leg. 26.2.2010 (10 shells; 2 spirit specimens, 1 dissected, MBC); Lourdes (department of Hautes-Pyrénées), 30TYN47, collector unknown (4 shells, ex Paulucci collection; FGC 19692). Spain, Comunidad Autónoma de Cantabria: Hazas-Asón (municipality of Santander), 30TV5282, B. Gómez leg. 26.1.1986 (5 spirit specimens, 3
Diagnosis

A genus of the aezcids characterised by ovoid-fusiform shell with callous rim on columella and parietum ending in transversely elongate angular tubercle, sinuous (more or less inverted S-like) outer margin with upper third thinner than elsewhere, composite apertural armature (consisting of at least five often knob/fold-like pieces: supraparietal, parietal, columellar lamellae and subcolumellar and upper palatal plicae), and protoconch smooth; radula without latero-marginal teeth; male distal genitalia with vas deferens almost uniform in diameter, penial retracttor inserted basally, muscular sheath on proximal penis, no penial papilla at opening of vas deferens into penis and simple longitudinal pleats and poorly discernible cordon inside penis.

Description of type species

Shell (Figs 10–13): dextral, small, imperforate, elongate ovoid-fusiform, reddish-brown, yellowish or colourless, glossy and transparent when fresh, with 7–8 almost flat whorls separated by superficial sutures; aperture orthocline, oblique pyriform, pointed at base, with apertural armature usually consisting of 7–8 parts (small supraparietal lamella; rather high oblique parietal lamella, sometimes anteriorly joined by transverse bridge to infraparietal lamella forming U-shaped structure; small to very small, sometimes absent, infraparietal lamella; descending columellar lamella; well evident subcolumellar lamella; one or two variably developed, immersed, lower palatal tooth-like plica; upper palatal tooth-like plica); peristome not reflected, but retracted with respect to last whorl (it seems to form a varix), slightly thickened in its outer upper third and greatly thickened elsewhere, with callous rim on columella and parietum sometimes continuous, evident and ending in transversely elongate angular tubercle separated by deep notch from upper vertex of aperture and with sinuous (more or less inverted S-like) outer margin (parietal lamella, angular tubercle and upper palatal knob delimit more or less evident sinus); protoconch smooth; teleoconch smooth with very thin spiral grooves. Shell dimensions: H – 6.4–8.0 mm; D – 2.7–2.8 mm.

Radula: central tooth small, tricuspid; lateral teeth large, bicuspid; latero-marginal teeth absent; marginal teeth pluricuspid (up to 12 cusps) (radular formula: M 16-17/5-12 + L 8-9/2 + C 1/3 + L 8-9/2 + M 16-17/5-12).

Female distal genitalia (Figs 14, 16–19, 23, 26): free oviduct very short to long and variably wide. Bursa copulatrix sac-like, bean-like or pyriform with long and slender to wide duct, initially not flared. Vagina short to long, variably wide, completely or partially (medium-distal portion) covered by usually faint, yellowish to brownish muff; internal surface smooth.

Male distal genitalia (Figs 14–28): vas deferens very long (longer than FO + V + P), almost uniform in diameter (just slightly thickened in its beginning

Figs 10–13. Shells of *Azeca goodalli* (Férussac, 1821): 10–12 – specimens from Lourdes (department of Hautes-Pyrénées), collector unknown; 13 – Eversden (county of Cambridgeshire), M. J. Bishop leg. 27.5.1972. Scale bar 2 mm
Figs 14–16. Genitalia of *Azeca goodalli* (Férussac, 1821): 14 – general view (gonad excluded); 15 – detail of penial complex; 16 – internal structure of distal penis; a specimen from Paythorne Bridge (county of Lancashire), D. LINDLEY & A. NORRIS leg. 13.4.2013. Scale bar 1 mm.

Figs 17–18. Genitalia of *Azeca goodalli* (Férussac, 1821): 17 – general view (gonad excluded); 18 – detail of penial complex and internal structure of distal penis; a specimen from Paythorne Bridge (county of Lancashire), D. LINDLEY & A. NORRIS leg. 13.4.2013. Scale bar 1 mm.
and subterminal section), entering penis basally or laterally; opening of vas deferens into penis simple (penial papilla absent). Penis rather long, subcylindrical, divided into distinct proximal and distal parts by slight constriction; proximal part with or without penial diverticulum and with penial sheath. Penial diverticulum, when present, very small to small, variable in shape and size (sac-like, elongate straight or hook-like). Penial sheath muf-like, enveloping final section of vas deferens and proximal penis, proximally usually continuous with penial retractor and ending distally level with slight constriction that divides proximal and distal portions of penis; sometimes part of proximal penis and its diverticulum (when present) protruding out of it. Penial retractor rather short and robust, usually inserted where the vas deferens enters the penis and usually continuous with penial sheath; sometimes thin branch of penial retractor joins apex of penial diverticulum. Internal surface of penis with very small simple longitudinal pleats in proximal portion and some (1–3) poorly visible, low, spongy cordons in distal portion.

Remarks

Azeca was introduced by Fleming (1828) for a species from the British Isles, Azeca tridens Fleming, 1828. It was subsequently regarded as a valid genus with different family allocations: in the sigmurethran family Ferussaciidae (Pilsbr 1908, Hesse 1922, in the subfamily Cochlicopinae) or in the orthurethran families Pupillidae (Watson 1920, in the subfamily Azecinae), Cochlicopidae (Thiele 1931, Zilch 1959, Schileyko 1998a) or Azecidae (Bank et al. 2001).

Phylogeny based on morphological characters (Fig. 6) shows that the monotypic Azeca, together with Hypnophila and the Gomphroa species except G. boissi, belongs to the main azecid clade based on the transversely elongate tubercle on the outermost parietum (3(1)); within the main azecid clade Azeca is the sister group of two Hypnophila species based on loss of rows of pits on the protoconch (14(0)).

Azeca is distinguished by a long series of shell and gross anatomical autapomorphies: supraparietal denticle/tubercle present (4(1)); parietal lamella present (5(1)); columellar lamella present (8(1)); upper palatal denticle/tubercle present (11(1)); outer peristomal margin in lateral view sinuous (inverted S-like) (13(1)); latero-marginal teeth absent (16(1)); muscular sheath on proximal penis present (23(1)).

Molecular phylogenies (Figs 7–9) show that Azeca is distant from all the other azecids but its relationships are variable and without support. Unfortunately no molecular data are available on Dalmatian Hypnophila and this prevents us from discussing its

Figs 19–22. Genitalia of Azeca goodalli (Férussac, 1821): 19, 21 – distal genitalia; 20, 22 – detail of penial complex and internal structure of distal penis; specimens from Louannec (department of Côte-D’Armor), M. Calcaño & S. Cianfanneli leg. 30.6.1993. Scale bar 1 mm
Figs 23–25. Genitalia of *Azeca goodalli* (Férussac, 1821): 23 – distal genitalia; 24–25 – detail of penial complex and internal structure of distal penis; specimens from Pic de Gillers (department of Pyrénées-Atlantiques), B. GÓMEZ leg. 30.10.1987 (23–24) and Isla (municipality of Santander), B. GÓMEZ leg. 26.12.1985 (25). Scale bar 1 mm.

Figs 26–28. Genitalia of *Azeca goodalli* (Férussac, 1821): 26 – distal genitalia; 27–28 – detail of penial complex and internal structure of distal penis; specimens from Hazas-Asón (municipality of Santander), B. GÓMEZ leg. 26.1.1986 (26–27) and Sainte-Engrâce (department of Pyrénées-Atlantiques), B. GÓMEZ leg. 28.7.1982 (28). Scale bar 1 mm.
relationships with the genus which are indicated by the phylogeny based on morphological characters.

The genus is monotypic and includes only the widespread *Azeca goodalli* (Férussac, 1821) which occurs in western Europe from Britain eastward to Germany and southward to the northern Iberian Peninsula (*Kerney et al. 1983, Welter-Schultes 2012*). The species is little known anatomically and some aspects remain uncertain and controversial (e.g. presence of epiphallus and structure of ovispermiduct): the original contributions are limited to *Boycott* (1919), *Hesse* (1922; partly based on Fritz Wiegmanns’ research), *Gómez* (1988) and *Gómez & Angulo* (1990). Early anatomical descriptions (*Boycott* 1919, *Hesse* 1922) overlooked the penial sheath, which when extended and intact, may conceal the presence of a penial diverticulum.

*Hesse* (1922) and *Gómez* (1988, 1991) claimed the existence of an epiphallus, however the difference in calibre between the proximal and distal sections of the duct connecting the prostatic ovispermiduct to the penis is minimal. It is therefore impossible to call it an epiphallus (a similar situation occurs in *Cryptazeca*).

*Gómez & Angulo* (1990) and *Gómez* (1991) stated that further features concern the ovispermiduct, namely a simple fertilisation pouch surrounded by subepithelial goblet gland cells (*Gómez* 1991: 99); an “oviductal caecum” in the proximal region of the female ovispermiduct (*Gómez & Angulo* 1990: 109, *Gómez* 1991: 99); a “blind-ending duct” in the proximal region of the ovispermiduct assumed to be homologous with the allospermiduct of the other stylommatophorans (*Gómez & Angulo* 1990: 109, 110). A fertilisation pouch with similar structure and a blind-ending duct of the ovispermiduct was also described in *Cryptazeca* (*Gómez* 1991: 96, 99). The blind-ending duct was considered by *Gómez & Angulo* (1990) to correspond to the diverticulum described by *Boycott* (1919). Our light microscope (Wild M5A) study of azedic anatomy failed to find the blind-ending diverticulum (likewise F. Wiegmanns and P. Hesse; *Hesse* 1922) or the oviduct caecum in any of the taxa examined.

The shell and anatomical variation of *Azeca goodalli* is remarkable. Shell variation mainly concerns the apertural armature and led to establishment of some variety-level taxa in the early literature (for example, see *Pilsbrý* 1908). Anatomical variations especially concern the proximal part of the penis, with at least five different patterns (*Table 5*). We do not know if this anatomical variation reflects taxonomic differentiation, artefacts due to pre-mortem stress, or different sexual maturation. However, the fact that some of these patterns coexist within a population suggests that this variation occurs at least partly within populations and does not support the differentiation of species.

*Table 5. Anatomical variation in Azeca goodalli populations mainly concerns proximal penis, of which there are at least five patterns*

|                  | Group 1 | Group 2 | Group 3 | Group 4 | Group 5 |
|------------------|---------|---------|---------|---------|---------|
| **penis**        | with diverticulum | with diverticulum | without diverticulum | with diverticulum | without diverticulum |
| **penial retractor** | branched: larger branch on distal vas deferens, smaller branch on apex of penial diverticulum | branched: larger branch continuous with penial sheath, smaller branch on apex of penial diverticulum | enveloping proximal penis, but often with lateral cleft from which part of proximal penis protrudes | enveloping proximal penis, but with lateral cleft from which part of proximal penis protrudes | enveloping proximal penis, but sometimes with lateral cleft from which part of proximal penis protrudes |
| **penial sheath** | enveloping part of proximal penis | enveloping proximal penis, but with lateral cleft from which part of proximal penis protrudes | enveloping part of proximal penis, including its diverticulum which is thus concealed | enveloping part of proximal penis, including its diverticulum which is thus concealed | enveloping part of proximal penis, including its diverticulum which is thus concealed |
| **populations**  | Spain: Alisas (*Gómez* 1988: pl. 3, fig. 3); Aránzasu (*Gómez* 1988: pl. 3, fig. 5); Hayal de Santiago (*Gómez* 1988: pl. 3, figs 1–2); ? Germany: Harz (*Hesse* 1922: pl. 1, fig. 7; pl. 2, fig. 8) | France: Louannec (*Figs 19–20, 21–22*); Spain: Isla (*Fig. 25*) | Spain: Hazas-Asón (*Fig. 26*) | Spain: Kakouetta (*Gómez* 1988: pl. 3, fig. 4); UK: Paythorne Bridge (*Figs 14–16*) | Spain: Hazas-Asón; France: Lourdes, Pic de Gilliers (*Figs 23–24*), Saint-Engrâce (*Fig. 28*); UK: Paythorne Bridge (*Figs 17–18*) |
Genus: Cryptazeca Folin et Bérillon, 1877(b)

Figs 29–41

Type species: Azeca monodonta Folin et Bérillon, 1877(a), by original designation

Material examined
Cryptazeca monodonta (Folin et Bérillon, 1877(a)) (Figs 29–37) – France, Nouvelle-Aquitaine: Eaux Bonnes (Pyrénées-Atlantiques), B. GÓMEZ leg. 28.10.1987 (4 shells, FGC 48677); Gorges de Kakuetta (Pyrénées-Atlantiques), B. GÓMEZ leg. 21.5.2013 (4 spirit specimens, 2 dissected, FGC 41610); Grottes de Sare (Pyrénées-Atlantiques), 30TXN1591, D. T. HOLYOAK & G. A. HOLYOAK leg. 7.11.2011 (1 spirit specimen dissected, CGAH).

Cryptazeca vasconica (Kobelt, 1894) (Figs 38–41) – Spain, Basque Country: W of Ranero (district of Vizcaya), 30TVN6990, D. T. HOLYOAK & G. A. HOLYOAK leg. 14–15.6.2007 (3 spirit specimens, 2 dissected, CGAH).

Diagnosis
A genus of the azecids characterised by conical/ovoid/slender-fusiform shell with sinuous (more or less inverted S-like) outer margin with upper third thinner than elsewhere, apertural armature only consisting of evident subcolumnellar lamella; peristome not reflected, thin in its outer upper third, slightly thickened elsewhere, continuous, with slightly sinuous (more or less inverted S-like) outer margin; protoconch with close, spiral rows of very small pits in grooves; teleoconch smooth, with or without weak collabral growth lines and with very thin spiral grooves. Shell dimensions: H – 3.1–3.8 mm; D – 1.5–1.7 mm.

Body: ocular spots present.

Radula: central tooth small, tricuspid; lateral teeth large, bicuspid; marginal teeth pluricuspid (Folin 1891: figs 1–3 bis) (radular formula unknown).

Female distal genitalia (Figs 33–36): free oviduct short and wide. Bursa copulatrix oval or pyriform with long slender duct, initially very flared. Vagina long and wide, medially covered by thick, brownish muff. Internally protuberance covered in small papillae with apical thorns (vaginal part of auxiliary copulatory organ, according to GÓMEZ 1991) where free oviduct and duct of bursa copulatrix join vagina.

Male distal genitalia (Figs 33–37): vas deferens rather long, uniform in diameter (thin for its entire length), entering penis medially on small protuberance (considered “a short and rudimentary epiphalus” by GÓMEZ 1991); opening of vas deferens into penis simple (penial papilla absent). Penis short, ovoid to pear-shaped, undivided, without penial sheath and penial diverticulum. Penial retractor very short and robust, inserted basally. Internal surface of penis with bulge completely covered in small papillae.
with apically hooked thorns (ring-shaped stimulatory organ according to GÓMEZ & ANGULO 1987 and GÓMEZ 1990b, c; ring-shaped sarcobelum according to GÓMEZ 1991).

Remarks

Cryptazeca was established by FOLIN & BÉRILLON (1877b) for a species from the Pyrénées-Atlantiques, France. It was regarded as a valid distinct genus in the sigmurethran families Ferussaciidae (e.g. PILSBRY 1908, HESSE 1922, GERMAIN 1930, THIELE 1931, ZILCH 1959, BANK et al. 2001, SCHILEYKO 1999, in the subfamily Cryptazecinae) or Subulinidae (GITTENBERGER 1983) by all subsequent authors. In the late 1980s, GÓMEZ & ANGULO (1987) showed that its type-species had an orthurethrous kidney, so they moved the genus to the orthurethran family Cochlicopidae. Subsequently MADEIRA et al. (2010) re-examined its relationships with the genera Azeca, Cochlicopa, Ferussacia and Hypnophila based on rRNA gene-cluster. They found that Cryptazeca belonged to a clade distinct from the one including Cochlicopa. Indeed Cryptazeca groups with Azeca and Hypnophila to form a clade which has unresolved relationships with the chondrinids and with a large assemblage including all the orthurethran genera examined. On the contrary Cochlicopa groups with the Hawaiian amastrid Leptachatina Gould, 1848, forming a clade nested in the large orthurethran assemblage, with sister group relationships to the valloniids but without statistical support.

Phylogeny based on morphological characters (Fig. 6) shows that Cryptazeca species constitute a supported clade characterised by at least three autapomorphies, namely sinuous outer peristomal margin in lateral view (13(1)), penial bulge (31(1)) completely covered in small papillae with apically hooked thorns (30(1)). In turn Cryptazeca has sister group relationships with the clade consisting of Gomezziella girottii, Gomphroa boissii and the main azecid clade.

Figs 33–37. Genitalia of Cryptazeca monodonta (De Folin et Bérillon, 1877(a)): 33, 36 – general view (gonad excluded); 34–35, 37 – internal structure of penis; specimens from Gorges de Kakuetta (Pyrénées-Atlantiques), B. GÓMEZ leg. 21.5.2013 (33–35) and Grottes de Sare (Pyrénées-Atlantiques), D. T. HOLYOAK & G. A. HOLYOAK leg. 7.11.2011 (36–37). Scale bar 1 mm
Supraspecific systematics of Azecidae

(Azeca goodalli and all the species currently assigned to Hypnophila), sharing at least two synapomorphies: upper third of outer peristome thinner than elsewhere (12(1)) and absence of a true penial papilla (28(3)).

The ML tree based on COI sequences (Fig. 7) suggests that Cryptazeca species form a clade separate from the Azeca plus Gomphroa group and from Hypnocarnica micaelae. On the contrary, the ML tree of ITS2 sequences (Fig. 8) shows that Cryptazeca is paraphyletic: Cryptazeca monodonta forms a clade with H. micaelae and the group has sister relationships with Gomphroa; in turn the larger group (C. monodonta, H. micaelae and Gomphroa) has sister group relationships with Cryptazeca speleae. What is more, in the ML tree of concatenated COI+ITS2 sequences (Fig. 9) C. monodonta clusters together with H. micae and in the larger group with C. spelaea forms a sister clade to Gomphroa.

As in Azeca, the presence of an epiphallus and the structure of the ovispermiduct remain unclear. The epiphallus is absent in C. monodonta and C. vasconica according to Gómez & Angulo (1987); a rudimentary epiphallus is present in C. monodonta according to Gómez (1991); a normal but only slightly developed epiphallus is present in C. spelaea and C. elongata (Gómez 1990b, c). However the calibre difference between the proximal and distal sections of the duct connecting the prostatic ovispermiduct to the penis is minimal. It is therefore impossible to call it an epiphallus. According to Gómez (1991), the ovispermiduct has a “blind-ending duct” that opens proximally into the free oviduct; this duct is assumed to be similar to that reported in Azeca and considered homologous to the allospermiduct of the other stylommatophorans by Gómez & Angulo (1990). However we failed to find it during our light microscope study (Wild M5A) of some specimens of Cryptazeca monodonta.

The penial bulge completely covered in small papillae with apically hooked thorns distinguishes Cryptazeca sharply from all the other azecid genera. However, homology based on shape and location may be postulated between the penial bulge of Cryptazeca species and the tongue-like structure in the proximal penis of Hypnocarnica micaelae.

Apart from the type species, Cryptazeca includes three other species: C. elongata Gómez, 1990(b), C. spelaea Gómez, 1990(c), and C. subcylindrica Folin et Béllin, 1877(b) (Holyoak & Holyoak 2012, Welter-Schultes 2012). However, the status of certain allopatric taxa synonymised by Holyoak & Holyoak (2012), based only on conchological characters, deserves further research (C. kobelti Gittenberger, 1983; C. vasconica (Kobelt, 1894)). Some species are known anatomically (C. elongata; C. monodonta, C. spelaea, C. vasconica; see: Gómez & Angulo 1987, Gómez1990a, b, 1991). Cryptazeca species differ anatomically by virtue of the insertion of the vas deferens into the penis (lateral on a small protuberance in C. monodonta; basal and simple in the other species), the vaginal muff (well developed in C. monodonta; only slightly developed in the other species), the vaginal part of the auxiliary copulatory organ (present in C. monodonta; absent in the other species?), the shape of the penial auxiliary copulatory organ (wider in C. monodonta; narrower in C. spelaea) and thorn shape (hooked in C. monodonta; not
hooked in *C. elongata* and *C. spelaea*); other differences concern vagina length and penis shape but it is difficult to state if they are true differences or due to different sexual maturation or artefacts of pre-mortem stress (see for example Gómez 1990b: figs 9 and 11).

**Genus: Gomeziella Cianfanelli, Bodon, Giusti et Manganelli, 2018(a)**

**Figs 42–49**

**Type species: Hypnophila girottii Esu, 1978**

**Material examined**

*Gomeziella girottii* (Esu, 1978) – Italy, Sardinia: Codula Fuili (province of Nuoro), 32TNK5256, S. CIANFANELLI & E. TALENTI leg. 19.5.2004 (126 shells, 3 spirit specimens, 2 dissected, MZUF GC/21905; 3 spirit specimens, 2 dissected, MZUF GC/21928); Punta Pedra Longa (province of Ogliastra), 32TNK6031, A. MARCELLI leg. 27.5.2013 and 21.6.2013 (93 shells, AMC; 2 specimens, 1 dissected, MZUF GC/43136, GC/43137); State Road 129, near San Leonardo, between Onifai and Orosei (province of Nuoro), 32TNK5671, S. CIANFANELLI & E. TALENTI leg. 27.10.1995 (85 shells, MZUF GC/42228; 4 shells, SCC).

**Diagnosis**

A genus of the azeceans characterised by very elongate, cylindrical to conical shell with sinusous (more or less inverted S-like) outer margin with upper third thinner than elsewhere, simplified apertural armature (consisting only of small, faint, often absent tubercle on innermost parietum and subcolumellar lamella) and protoconch with rows of pits; radula with latero-marginal teeth; male distal genitalia with vas deferens thickened submedially, penial retractor inserted laterally, no muscular sheath on proximal penis, no penial papilla at opening of vas deferens into penis and longitudinal pleats, one crest-like, inside penis.

**Description of type species**

**Shell (Figs 42–46):** dextral, small, imperforate, elongate to very elongate, cylindrical-conical or cylindro-fusiform, colourless, glossy and transparent when fresh, with 7½–10 slightly convex to flat whorls, separated by superficial sutures; aperture slightly prosocline, oblique pyriform, rounded at base, with simplified apertural armature (consisting of only small, faint, often absent, tubercle on innermost parietum and barely evident subcolumellar lamella); peristome not reflected, thin in outer upper third, slightly thickened elsewhere (point of passage sometimes knob-like) and more or less continuous, with sinusous (more or less inverted S-like) outer margin; protoconch with close, spiral rows of very small pits in grooves; teleoconch smooth, with very thin spiral grooves. Shell dimensions: H – 4.0–6.8 mm; D – 1.2–1.9 mm.

**Body:** ocular spots absent.

**Radula:** central tooth small, tricuspid; lateral teeth large, bicuspid; latero-marginal tooth tricuspid (because ectocone split into two); marginal teeth pluricuspid (up to 9 cusps) (radular formula: M ca. 10/4-9 + LM 1/3-4 + L 3/2 + C 1/3 + L 3/2 + LM 1/3-4 + M ca. 10/4-9).

**Female distal genitalia (Figs 47–49):** free oviduct short to long and variably wide. Bursa copulatrix...
bean-like or pyriform with long to very long, very slender duct, initially barely flared. Vagina long and rather slender, proximally covered by faint yellowish muff; internal surface smooth.

**Male distal genitalia** (Figs 47–49): vas deferens very long (longer than FO + V + P), uneven in diameter (thin for entire length except for submedial section which is markedly thickened) entering penis basally (subterminal section not enveloped by penial retractor); opening of vas deferens into penis simple (penial papilla absent). Penis long, subcylindrical (wider medio-distally and tapering distally), undivided, without penial sheath or penial diverticulum. Penial retractor short and robust, inserted laterally at about first third of penis length. Internal surface of proximal penis with two longitudinal pleats: one pleat with jagged edges continues uninterrupted for entire length of penis; the other pleat sometimes branches in two level with point of insertion of penial retractor; these two pleats continue for part or entire length of distal penis (one of the latter two is sometimes higher, sinuuous and crest-like).

**Remarks**

*Gomeziella girottii* does not belong to the ferus-sacids, contrary to the claim of **Welter-Schultes** (2012): its orthurethrous kidney definitively excludes this possibility.

Phylogeny based on morphological characters (Fig. 6) shows that the species is characterised by at least five autapomorphies, namely very elongate, cylindrical-conical or cylindrical-fusiform shell (1(3)), sinuous outer peristomal margin in lateral view (13(1)), eye spots absent (15(1)), vas deferens well thickened submedially (19(1)) and lateral insertion of penial retractor (27(0)); in turn *G. girottii* has unresolved relationships with *Gomphroa boissii* and the main azecid clade including *Azeca* and all the species currently assigned to *Hypnophila*. This group (*Gomeziella girottii*, *Gomphroa boissii* and the main azecid clade) shares at least one synapomorphy: the penial longitudinal pleats (29(1)).

One of the most peculiar features of *G. girottii* is a submedially well thickened vas deferens. A similar vas deferens structure is known in few other unrelat-
ed taxa (e.g. orculids: *Alvariella* Hausdorf, 1996 and *Sphyradium* Charpentier, 1837; see Hausdorf 1996).

The new genus includes only one species from Sardinia (western Mediterranean). Its populations vary in shell shape and size, but it is still unclear whether this variation reflects taxonomic differentiation.

**Genus: Gomphroa** Westerlund, 1902

**Figs 50–80**

**Type species:** *Zua boissii* Dupuy, 1851, by monotypy

**Material examined**

*Gomphroa boissii* (Dupuy, 1851) (Figs 50–54) – France, Provence-Alpes-Côte-d’Azur: Callelongue (department of Bouches-du-Rhône), 31TFH9186, M. Bodon leg. 30.12.1989 (7 shells, MBC); Spain, Cataluña: Collserola, Ermita de Sant Vicenç (province of Barcelona), V. Bros leg. 8.2002 (2 spirit specimens dissected; FGC 41611); Las Planas de Vallvidrera (province of Barcelona), C. ALTIMIRA leg. 4.1956 (3 shells, FGC 16563); Les Teixoneres (province of Barcelona), J. COMAS NAVARRO leg. 9.9.2002 (2 shells, MZB 2009-0646); Parc Natural de Sant Llorenc del Munt (province of Barcelona), V. Bros leg. 2.2013 (1 spirit specimen dissected, FGC 41615); Queralt (province of Barcelona), G. GARDINI leg. 22.7.1990 (1 spirit specimen dissected, FGC 41617).

*Gomphroa dohrni* (Paulucci, 1882) (Figs 55–58) – Italy, Sardinia: Costa Dorata (province of Olbia-Tempio Pausania), 32TNL5524, S. CIANFANELLI & E. Lori leg. 1.5.2003 (2 spirit specimens, 1 dissected, MZUF GC/7573); Monte Nieddu (province of Carbonia-Iglesias), collector unknown leg. 23.3.1976 (1 spirit specimen dissected, FGC 29598); Orroli (province of Cagliari), 32SNJ29, L. PINTER leg. 2.4.1978 (3 spirit specimens, 1 dissected, FGC 29596).

*Gomphroa cf. remyi* (Boettger, 1949) (Figs 59–61) – France, Corsica: Cap Corse, Camera, 32TNNN2956, S. TATTI leg. 18.4.1981 (1 spirit specimen, FGC 29615); Cap Corse, Pietra Nera, 32TNNN3730, L. CASTAÑOLO, F. GIUSTI & G. MANGANELLI leg. 4.12.1983 (3 spirit specimens, 2 dissected, FGC 29624).

*Gomphroa bisacchii* (Giusti, 1970) (Figs 62–65) – Italy, Sardinia: Cala Gonone (province of Nuoro), 32TNK55, F. GIUSTI leg. 2.5.1969 (1 shell [holotype], 2 spirit specimens [paratypes] dissected, FGC 9290, 19648); Cala Gonone, Codula Fuili (province of Nuoro), 32TNK5356, S. CIANFANELLI & E. TALENTI leg. 19.5.2004 (6 spirit specimens, 2 dissected, MZUF GC/21929); Dorgali (province of Nuoro), 500–600 m a.s.l., 32TNK56, L. PINTER leg. 5.4.1978 (7 shells, FGC 16561; 9 spirit specimens, 2 dissected, FGC 29591); Dorgali (province of Nuoro), 32TNK56, collector unknown leg. 22.5.1980 (11 spirit specimens, 2 dissected, FGC 29590).

*Gomphroa etrusca* (Paulucci, 1886) (Figs 66–68) – Italy, Tuscany, Monte Argentario (province of Grosseto): Porto Santo Stefano, not georeferenceable, collector and date unknown (1 spirit specimen, 1 dissected, FGC 16908); cemetery of Porto Santo Stefano, 32TPM7499, V. SBDONI leg. 29.4.1978 (1 spirit specimen dissected, FGC 16909).

*Gomphroa cf. etrusca* (Paulucci, 1886) (Figs 69–72) – Italy, Tuscany, Tuscan Archipelago: Island of Giglio (province of Grosseto), Il Franco, 32TPMS9, L.

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Figs 50–53. Shells of *Gomphroa boissii* (Dupuy, 1851), specimens from Les Teixoneres (province of Barcelona), J. COMAS NAVARRO leg. 9.9.2002 (50–51) and Las Planas de Vallvidrera (province of Barcelona), C. ALTIMIRA leg. 4.1956 (52–53). Scale bar 2 mm
FAVILLI & G. MANGANELLI leg. 21.2.1990 (10 shells, FGC 5015), A. Benocci, F. GIUSTI & G. MANGANELLI leg. 27.2.2013 (6 shells, 2 spirit specimens dissected, FGC 40920); Island of Gorgona (province of Livorno), Torre Vecchia, F. Giusti leg. 31.3.1974 (2 spirit specimens dissected, FGC 16899).

Figs 54–55. Genitalia (gonad excluded) of: 54 – *Gomphroa boissii* (Dupuy, 1851), specimens from Collserola, Ermita de Sant Vicenç (province of Barcelona), V. Bros leg. 8.2002; 55 – *Gomphroa dohrni* (Paulucci, 1882) from Costa Dorata (province of Olbia-Tempio Pausania), S. Cianfanelli & E. Lori leg. 1.5.2003. Scale bar 1 mm

Figs 56–58. Genitalia of *Gomphroa dohrni* (Paulucci, 1882): 56–57 – general view (gonad excluded); 58 – internal structure of penis; specimens from Orroli (province of Cagliari), L. Pintér leg. 2.4.1978 (56) and Monte Nieddu (province of Carbonia-Iglesias), collector unknown, 23.3.1976 (57–58). Scale bar 1 mm
Figs 59–61. Genitalia of Gomphroa cf. remyi (Boettger, 1949): 59–60 – general view (gonad excluded); 61 – internal structure of penis; specimens from Cap Corse, Pietra Nera, L. Castagnolo, F. Giusti & G. Manganelli leg. 4.12.1983. Scale bar 1 mm

Figs 62–65. Genitalia of Gomphroa bisacchii (Giusti, 1970): 62 – general view (gonad excluded); 63–65 – internal structure of penis; specimens from Cala Gonone, Codula Fuili (province of Nuoro), S. Cianfanelli & E. Tarenti leg. 19.5.2004 (62–63, 65) and Cala Gonone (province of Nuoro), F. Giusti leg. 2.5.1969 (64). Scale bar 1 mm
Figs 66–68. Genitalia of *Gomphroa etrusca* (Paulucci, 1886): 66, 68 – general view (gonad excluded); 67 – internal structure of penis; specimens from Monte Argentario, cemetery of Porto Santo Stefano (province of Grosseto), V. S Bordoni leg. 29.4.1978 (66–67) and Porto Santo Stefano (province of Grosseto), collector and date unknown (68). Scale bar 1 mm.

Figs 69–70. Genitalia of *Gomphroa cf. etrusca* (Paulucci, 1886): 69 – general view (gonad excluded); 70 – internal structure of penis; a specimen from Island of Giglio, Il Franco (province of Grosseto), A. Benocci, F. Giusti & G. Manganelli leg. 27.2.2013. Scale bar 1 mm.
Figs 71–72. Genitalia of *Gomphroa cf. etrusca* (Paulucci, 1886): 71 – general view (gonad excluded); 72 – internal structure of penis; a specimen from Island of Gorgona, Torre Vecchia (province of Livorno), F. GIUSTI leg. 31.3.1974. Scale bar 1 mm

Figs 73–74. Genitalia of *Gomphroa incerta* (Bourguignat, 1858): 73 – general view (gonad excluded); 74 – internal structure of penis; a specimen from Island of Stromboli, Ginostra (province of Messina), F. GIUSTI leg. 3.11.1969. Scale bar 1 mm
Figs 75–76. Genitalia of Gomphroa cf. cylindracea (Bourguignat, 1858): 75 – general view (gonad excluded); 76 – internal structure of penis; a specimen from Santuario di Santa Maria della Scala (province of Siracusa), D. CARUSO leg. 13.10.1977. Scale bar 1 mm

Figs 77–78. Genitalia of Gomphroa cf. cylindracea (Bourguignat, 1858): 77 – general view (gonad excluded); 78 – internal structure of penis; a specimen from Santuario di Santa Maria della Scala (province of Siracusa), D. CARUSO leg. 13.10.1977. Scale bar 1 mm
Gomphroa incerta (Bourguignat, 1858) (Figs 73–74) – Italy, Sicily, Aeolian Islands: Island of Lipari, along the road to Monte Sant’Angelo (province of Messina), not georeferenceable, F. Giusti leg. 25.7.1972 (1 spirit specimen dissected, FGC 17277); Island of Panarea, slopes of Monte Alto (province of Messina), 33WC07, F. Giusti leg. 30.3.1971 (4 shells, 10 spirit specimens, 2 dissected, FGC 17273); Island of Stromboli, Ginostra, 33SWC1693, F. Giusti leg. 3.11.1969 (5 shells, 2 spirit specimens dissected, FGC 11005).

Gomphroa cf. cylindracea (Calcara, 1840) (Figs 75–78) – Sicily, Santuario di Santa Maria della Scala (province of Siracusa), 33SWA0291, D. Caruso leg. 13.10.1977 (4 spirit specimens, 2 dissected, FGC 41618).

Gomphroa emiliana (Bourguignat, 1858) (Figs 79–80) – Italy, Sicily, Egadi Islands: Island of Marettimo, above the village (province of Trapani), 33STC4206, F. Giusti leg. 2.4.1982 (6 shells, 29 spirit specimens, 4 dissected, FGC 17791).

Gomphroa zirjensis (Štamol, Manganelli, Barbato et Giusti, 2018) – Croatia, Šibenik Archipelago, Island of Žirje, Gradina (area), Gradina (pit), 33TWJ53, B. Jalžić leg. 10.2004 (31 shells, CNHM 10919), B. Jalžić leg. 10.2007 (7 specimens, 3 dissected, CNHM 10919).

Diagnosis

A genus of the azeicids characterised by ovoid-fusiform to ovoid-cylindrical shell with callous rim on columella and parietum ending in transversely elongate angular tubercle (absent in the type species), straight outer margin with upper third thinner than elsewhere, without apertural armature (apart from barely evident subcolumellar lamella) and having a protoconch with rows of pits; radula with latero-marginal teeth; male distal genitalia with vas deferens almost uniform in diameter, penial retractor inserted basally/sub-basally, no muscular sheath on proximal penis, penial papilla consisting of bunch of short conical digit-like appendages fused at base (sometimes forming cylindrical structure with fringed tip) at opening of vas deferens inside penis (presumably absent in the type species) and longitudinal pleats with knobs and transverse crests inside penis.
**Description of type species**

**Shell** (Figs 50–53): dextral, small, imperforate, elongate cylindrical-fusiform, yellowish, glossy and transparent when fresh, with 5–6 slightly convex to flat whorls, separated by superficial sutures; aperture slightly prosocline, ovate pyriform, rounded at base, without apertural armature (apart from barely evident subcolumellar lamella); peristome not reflected, not thickened (very thin in outer upper third), with callous rim on columella and parietum interrupted and very faint (transversely elongate angular tubercle absent) and with straight outer margin; protoconch with close spiral rows of very small pits in grooves; teleoconch smooth, with very thin spiral grooves. Shell dimensions: H – 4.9–5.8 mm; D – 1.7–2.1 mm.

**Body**: ocular spots present.

**Radula**: central tooth small, tricuspid; lateral teeth large, bicuspid; latero-marginal teeth tricuspid (because ectocone split into two); marginal teeth pluricuspid (radular formula unknown).

**Female distal genitalia** (Fig. 54): free oviduct long and wide. Bursa copulatrix oval or bean-like with short slender duct, initially barely flared. Vagina long and rather slender, almost completely covered by faint, pale yellow to brownish muff; internal surface smooth.

**Male distal genitalia** (Fig. 54): vas deferens rather long, almost uniform in diameter (slightly thickened proximally, then uniformly slender), entering penis basally (its final section not enveloped by penial retractor); opening of vas deferens into penis apparently simple. Penis very short, subcylindrical, undivided, without penial sheath or penial diverticulum. Penial retractor short and thin, inserted basally close to vas deferens. Internal surface of penis not clearly resolved (due to very small size of penis), apparently with two, thin, longitudinal pleats.

**Remarks**

*Gomphroa* was introduced as a subgenus of *Zua* by *Westerlund* (1902) for a species from the eastern Pyrenees. Later it was considered a subgenus of *Azeca* (e.g. *Pilsbry* 1908, *Hesse* 1922, *Germain* 1930, *Thiele* 1931) or a distinct genus (*Zilch* 1959, *Schileyko* 1998a) in the orthurethran family Cochlicopidae (*Zilch* 1959, *Schileyko* 1998a) or it was disregarded and its type species included in *Hypnophila* (*Giusti* 1976, *Gittenberger* 1983, *Gómez* 1990a, *Bank* et al. 2001, *Welter-Schultes* 2012).

Phylogeny based on morphological characters (Fig. 6) shows that *Gomphroa boissii* is characterised by at least one autapomorphy, namely very small penis (22(2)), and it has unresolved relationships with *Gomeziella girottii* and the main azecid clade including *Azeca* and all the species currently assigned to *Hypnophila*. This group (*Gomeziella girottii, Gomphroa boissii* and the main azecid clade) shares at least one synapomorphy: penial longitudinal pleats (29(1)). The species currently assigned to *Hypnophila* fall into two distinct lineages: one, including the Dalmatian species except *H. zirjensis*, is characterised by cup-like initial portion of penial plica (28(1)), and is the sister group of *Azeca goodalli* based on the absence of rows of pits on the protoconch (14(0)); the other, including all the western Mediterranean species plus the Dalmatian *H. zirjensis*, is characterised by an apically fringed papilla through which the vas deferens opens into the penis (28(2)). In the western Mediterranean *Hypnophila* lineage, *H. bisacchii* is sister group to all the other species examined, due to its elongate ovoid-fusiform shell (1(0)) (elongate cylindrical-fusiform shell in the others (1(2))); in this subclade, two species – *Hypnophila dohrni* and *H. reymyi* – have sister group relationships based on small penial size (22(1)).

The ML tree based on COI sequences (Fig. 7) shows that the western *Hypnophila* belong to two lineages: the first is well supported and includes species from southern France (*Hypnophila* sp. A), Sardinia (*H. bisacchii*), Tuscany and the Tuscan Archipelago (*H. etrusca*) and Pontine Archipelago (*Hypnophila* sp. B); the other is weakly supported and includes species from the Iberian Peninsula (*H. boissii* and *H. malagana*) and Sardinia and Corsica (*H. dohrni* and *H. reymyi*). The ML trees of ITS2 (Fig. 8) and of concatenated COI+ITS2 (Fig. 9) sequences show all these species as members of the same well supported lineage with three subclades: one including southern French, Sardinian and Tyrrenhian species (*Hypnophila* sp. A, *Hypnophila* sp. B, *H. bisacchii* and *H. etrusca*), one including Iberian species (*H. boissii* and *H. malagana*) and one including Sardinian and Corsican species (*H. dohrni* and *H. reymyi*).

It is evident that morphological and molecular phylogenetic analyses diverge regarding the relationships of *H. boissii*. Although we always emphasise that the interpretation of anatomical and molecular features should be consistent without prioritizing either of them, we think that in the light of molecular results a new interpretation of morphological data is possible. Therefore, we assign the western Mediterranean species to *Gomphroa*. After the present revision, most *Hypnophila* species come to fall in *Gomphroa*. Apart from the two species from North Africa (*G. psathyrolena* (Bourguignat, 1864) and *G. maroccana* (Mousson, 1873)), all the others are known anatomically, although some were studied for the first time here (*Giusti* 1968, 1970, 1973, 1976, *Gómez* 1990a, present paper).

*Gomphroa* species usually have an evident angular tubercle, a normally developed penis and a vas deferens which is bordered by a penial papilla consisting of a bunch of short conical digit-like appendages.
fused at the base (sometimes forming a cylindrical structure with fringed tip) where it opens into the penis.

Many *Gomphroa* species are very little known and are distinguished mainly on a geographical basis. Conchologically, only two species are very distinctive, the Sardinian *G. bisacchii* and the Iberian *G. malagana* (ŠtAMOL et al. 2018). Anatomical differences consist in the shape and size of the different sections of the distal genitalia. The Sardinian *G. dohrni* (Figs 55–58) and the Corsican *G. cf. remyi* (Figs 59–61) have a proportionally smaller penis; the Sicilian *G. cf. cylindracea* has a long penial papilla (Figs 76, 78; but see also the Tuscan *G. cf. etrusca* from Gorgona: Fig. 72); the Aeolian *G. incerta* has sub-basal insertion of the penial retractor (Fig. 73; but see also the Sicilian *G. cf. cylindracea* from Santuario di Santa Maria della Scala: Figs 75, 77, and the Tuscan *G. cf. etrusca* from Gorgona: Fig. 71). However, since very few specimens have been studied, it is difficult to evaluate the significance of these differences or to know whether they are due, for example, to species-specific divergence, different sexual maturation, different fixation or an artefact of pre-mortem stress. For example, this is the case of *G. etrusca*: specimens from some Tuscan islands (Giglio and Gorgona) differ slightly from those collected on Monte Argentario. Future molecular research is needed to verify the status at species level of the populations present in Italy and especially the single islands of the Tuscan Archipelago.

**Genus: Hypnocarnica** Cianfanelli et Bodon in Cianfanelli et al., 2018(b)

**Figs 81–89**

**Type species:** *Hypnocarnica micaelea* Cianfanelli et Bodon in Cianfanelli et al., 2018, by original designation

**Material examined**

*Hypnocarnica micaelea* Cianfanelli et Bodon in Cianfanelli et al., 2018 – Italy, Friuli – Venezia Giulia: southern slope of Monte Jouv (municipality of Maniago, province of Pordenone), 32TUM2015, M. Bodon & S. Cianfanelli leg. 20.8.2015 (5 shells, MZUF GC/47777; 3 shells, MZUF GC/47778), M. Calcagno & S. Cianfanelli leg. 20.4.2016 (91 shells, 1 specimen dissected, MZUF GC/50063); south-western slope of Monte Jouv (municipality of Maniago, province of Pordenone), 32TUM1916, E. Bodon, M. Bodon & S. Cianfanelli leg. 3.4.2015 (1 shell, MZUF GC/46982).

**Diagnosis**

A genus of the orthurethrans characterised by very elongate, cylindrical to conical shell with sinuous (more or less inverted S-like), thin outer margin, without apertural armature (apart from barely evident subcolumellar lamella) and protoconch with rows of pits; rudula with latero-marginal teeth; male distal genitalia with vas deferens almost uniform in diameter, penial retractor inserted basally, no muscular sheath on proximal penis, true penial papilla at opening of vas deferens into penis and large tongue-like structure, possibly a sort of sarcobelum or stimulatory organ inside final part of proximal penis.

**Description of type species**

**Shell** (Figs 81–86): dextral, small, imperforate, elongate to very elongate, cylindrical or cylindro-conical, colourless, glossy and transparent when fresh, with 5½–5¾ slightly convex to flat whorls, separated by superficial sutures; aperture slightly prosocline, oblique pyriform, rounded at base, with simplified apertural armature (consisting of barely evident subcolumellar lamella); peristome not reflected, thin along entire outer margin and more or less continuous, with sinuous (more or less inverted S-like) outer margin; protoconch smooth; teleconch with very thin spiral grooves particularly evident on last whorl near aperture. Shell dimensions: H = 4.4–8.3 mm; D = 1.3–2.3 mm.

**Body:** eye spots absent.

**Radula:** central tooth small, tricuspid; lateral teeth large, bicuspid; latero-marginal tooth tricuspid (ectocone split into two); marginal teeth pluricuspid (up to 9 cusps) (radular formula: M ca. 10/6-14 + LM 1/3-5 + L 4/2 + C 1/3 + L 4/2 + LM 1/3-5 + M ca. 10/6-14).

**Female distal genitalia** (Figs 87–88): free oviduct short. Bursa copulatrix oval with very long wide duct, initially not flared and rather slender at its end. Vagina short and wide, covered by thin brownish muf; internal surface smooth.

**Male distal genitalia** (Figs 87–89): vas deferens rather long, almost uniform in diameter, entering penis basally; opening of vas deferens into penis through true penial papilla. Penis long, divided into proximal and distal parts of different diameter; proximal part bean-like, without penial sheath or penial diverticulum, divided by slight constriction into small initial part internally with penial papilla and large final part internally with large tongue-like structure, possibly a sort of sarcobelum or stimulatory organ; distal part initially slender, widening distally. Penial retractor rather short and robust, inserted basally, close to vas deferens. Internal surface of penis smooth.
Figs 81–86. Shells of *Hypnocarnica micaelae* Cianfanelli et Bodon in Cianfanelli et al., 2018(b): 81 – a specimen from the south-western slope of Monte Jouf (province of Pordenone), E. BODON, M. BODON & S. CIANFANELLI leg. 3.4.2015; 82–85 – specimens from the southern slope of Monte Jouf (province of Pordenone), M. BODON & S. CIANFANELLI leg. 20.8.2015 (82 – MZUF 47778), M. BODON & S. CIANFANELLI leg. 20.8.2015 (83, 85 – MZUF 47777) and M. CALCAGNO & S. CIANFANELLI leg. 20.4.2016 (84, 86 – MZUF 50063). Scale bar 2 mm

Figs 87–89. Genitalia of *Hypnocarnica micaelae* Cianfanelli et Bodon in Cianfanelli et al., 2018(b): 87–88 – general view (gonad excluded); 89 – internal structure of proximal penis with penial papilla and tongue-like structure; a specimen from the southern slope of Monte Jouf (province of Pordenone), M. CALCAGNO & S. CIANFANELLI leg. 20.4.2016. Scale bar 1 mm
Remarks

Hypnocarnia was introduced for a new species from the Carnic Prealps, north-eastern Italy, assigned to the azecids on the basis of morphological and molecular characters. Its shell is very similar in shape, size and variation to that of species of Cecilioides with very slender shells, like those of C. acicula, and less slender shells like those of C. petitianus/tumulorum/janii. It is surprising that no name established for Cecilioides was available for this species. Although its shell is similar to that of Cecilioides, the orthurethran kidney excludes any relationship with the sigmurethran ferussacids.

Regarding the internal structure of the penis, Cianfanelli et al. (2018b) failed to detect the true penial papilla at the opening of vas deferens inside the proximal penis and interpreted the more distal tongue-like structure as a penial papilla.

Phylogeny based on morphological characters (Fig. 6) shows that H. micaele is the sister group of all the other azecids and is characterised by at least three autapomorphies, namely very elongate, cyindrical-conical or cyindrical-fusiform shell (1(4)), eye spots absent (15(1)) and tongue-like structure inside proximal penis (31(1)).

The Bayesian Inference tree based on concatenat-ed sequences by Cianfanelli et al. (2018b: fig. 7) shows that H. micaele is the sister group of all the other azecids and is characterised by at least three autapomorphies, namely very elongate, cylind-rical-conical or cyindrical-fusiform shell (1(4)), eye spots absent (15(1)) and tongue-like structure inside proximal penis (31(1)).

The type species Hypnophila zacynthia (Roth, 1855) – Greece, Iónia (Cantraine, 1835) – Croatia: Figs 93–95 (19505); Island of Kefalloniá, Póros, W. Rähle leg. 28.9.1980 (1 spirit specimen dissected, FG 17280); Island of Itháki, 0.8 km N of Itháki village, collector and date unknown (2 shells, FG 41742).

Diagnosis

A genus of the azecids characterised by ovoid-fusiform to ovoid-cylindrical shell with callous rim on columella and parietum ending in transversely elongate angular tubercle, almost straight outer margin with upper third thinner than elsewhere, without apertural armature (apart from barely evident subcolumellar lamella) and protoconch without rows of pits; radula with latero-marginal teeth; male distal genitalia with vas deferens almost uniform in diameter, penial retractor inserted basally, no muscular sheath on proximal penis and two longitudinal pleats with thickened parts that fuse proximally, giving rise to small pointed cup-like crest bordering opening of vas deferens inside penis.

Description of type species

Shell (Figs 90–92): dextral, small, imperforate, elongate ovoid-fusiform, pale-brownish or yellowish, glossy and transparent when fresh, with 6½–7, slightly convex to flat whorls, separated by superficial sutures; aperture slightly prosocline, ovate pyriform, rounded at base, without apertural armature (apart from barely evident subcolumellar lamella); peristome not reflected, not thickened (thin in its outer upper third), with callous rim on columella and parietum sometimes continuous, faint and ending in transversely elongate angular tubercle separated by deep notch from upper angle of aperture and with straight outer margin; protoconch with slightly raised, interrupted radial crests; teleoconch smooth, with very thin spiral grooves. Shell dimensions: H – 6.5–7.6 mm; D – 2.6–3.3 mm according to Gittenberger (1993).

Body: eye spots present.

Radula: central tooth small, tricuspid; lateral teeth large, bicuspid; latero-marginal teeth tricuspid (because ectocone split into two); marginal teeth pluri-cuspid (up to 9 cusps) (radular formula: M 13/3–9 + LM 2/3 + L 8/2 + C 1/3 + L 8/2 + LM 2/3 + M 13/3–9).

Female distal genitalia (Fig. 93): free oviduct long and wide. Bursa copulatrix oval or pyriform with long and rather wide duct, initially barely flared. Vagina short and wide, proximally covered by thick, brownish muf; internal surface smooth.

Male distal genitalia (Figs 93–95): vas deferens rather long, almost uniform in diameter (slightly thickened proximally, then uniformly slender), entering penis basally (its final section more or less enveloped by penial retractor); opening of vas deferens

Genus: Hypnophila Bourguignat, 1858

Figs 90–95

Type species: Bulimus pupaformis Cantraine, 1835, by subsequent designation (Pilsbry 1908)

Material examined

Hypnophila pupaformis (Cantraine, 1835) – Croatia: Dubrovnik, collector unknown leg. 9.1969 (3 shells, FG 19589); Island of Vis, Komiza, F. Giusti leg. 23.7.1968 (5 shells, FG 19009); Sipun pećina, Cavtat, E. Gittenberger leg. 2.5.1974 (1 spirit specimen dissected, FG 19010); Sipun pećina, Cavtat, R.O. Izevedely leg. 7.5.2012 (3 spirit specimens, 1 dissected, FG 41613), B. Jaličič 15.5.2018 (8 spirit specimens, 2 dissected, FG 48643).

Hypnophila zacynthia (Roth, 1855) – Greece, Iónia. Nísia: Island of Kefalloniá, Karavomilos, E. Gittenberger leg. 6.5.1979 (2 shells, FG 19505).
Supraspecific systematics of Azecidae

Figs 90–92. Shells of *Hypnophila pupaeformis* (Cantraine, 1835): specimens from Island of Vis, Komiža, F. GIUSTI leg. 23.7.1968. Scale bar 2 mm.

Figs 93–95. Genitalia of *Hypnophila pupaeformis* (Cantraine, 1835): 93 – general view (gonad excluded); 94–95 – internal structure of penis; specimens from Sipun pećina at Cavtat, E. GITTENBERGER leg. 2.5.1974 (93–94) and R.O. IZVEDELY leg. 7.5.2012 (95). Scale bar 1 mm.
into penis bordered by initial section of shorter penial pleat. Penis long, uniformly cylindrical, undivided, without penial sheath or penial diverticulum. Penial retractor rather long and robust, inserted basally, enveloping final section of vas deferens. Internal surface of penis with two longitudinal pleats that arise proximally; shorter pleat runs for 2/3 of penis length and at its beginning gives rise to small point-ed cup-like crest bordering opening of vas deferens into penis; longer pleat, initially very slender then progressively larger, runs for entire length of penis and ends close to genital atrium, fringing or giving rise to thickened parts.

Remarks

Hypnophila was established for a “series” of Azeca by Bourguignat (1858) with six species originally included. It was subsequently regarded as a subgenus of Azeca (Pilsbry 1908, Hesse 1922, Germain 1930, Thiele 1931) or a distinct genus (Zilch 1959, Bank et al. 2001) in the orthurethran families Cochlicopidae (Zilch 1959) or Azeceans (Bank et al. 2001).

Phylogeny based on morphological characters (Fig. 6) shows that Hypnophila species belong to a monophyletic group supported by two synapomorphies: the elongate ovoid-cylindrical shell (1(2)) and the cup-like initial portion of one of the two penial pli-cae bordering the vas deferens opening into the penis (28(1)). This clade constitutes the sister group of Azeca based on loss of the rows of pits on protoconch (14(0)); in turn, Azeca plus Hypnophila are the sister group of the lineage including Gomphra species except G. boissii based on the transversely elongate tubercle on the outermost parietum (3(1)). Unfortunately no molecular data are available on Hypnophila and this prevents any discussion of its relationships revealed by phylogeny based on morphological characters.

Currently Hypnophila includes 15 species (Giusti & Manganelli 1984, Welter-Schultes 2012, Štamoš et al. 2018) but after the present revision, only four remain, all from western Balkan Peninsula and from western Greece, islands included: the type species (H. pupaeformis) plus H. polita (Porro, 1838), H. cyclothya (Boettger, 1885) and H. zacynthia (Roth, 1855) (GittenberGer 1993). These three species differ from H. pupaeformis and from each other in very few shell characters and in some cases show overlapping distributions (GittenberGer 1993: fig. 5).

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