Three redescriptions in Tintinnopsis (Protista: Ciliophora: Tintinnina) from coastal waters of China, with cytology and phylogenetic analyses based on ribosomal RNA genes

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Abstract

**Background:** The taxonomy of tintinnine ciliates is vastly unresolved because it has traditionally been based on the lorica (a secreted shell) and it has only recently incorporated cytological and molecular information. *Tintinnopsis*, the most speciose tintinnine genus, is also the most problematic: it is known to be non-monophyletic, but it cannot be revised until more of its species are studied with modern methods.

**Results:** Here, *T. hemispiralis* Yin, 1956, *T. kiaochowensis* Yin, 1956, and *T. uruguayensis* Balech, 1948, from coastal waters of China, were studied. Lorica and cell features were morphometrically investigated in living and protargol-stained specimens, and sequences of three ribosomal RNA (rRNA) loci were phylogenetically analyzed. The three species show a complex ciliary pattern (with ventral, dorsal, and posterior kineties and right, left, and lateral ciliary fields), but differ in lorica morphology, details of the somatic ciliature and rRNA gene sequences. *Tintinnopsis hemispiralis* is further distinguished by a ciliary tuft (a ribbon of very long cilia originated from the middle portion of the ventral kinety and extending out of the lorica) and multiple macronuclear nodules. Both *T. kiaochowensis* and *T. uruguayensis* have two macronuclear nodules, but differ in the number of somatic kineties and the position of the posterior kinety. Two neotypes are fixed for *T. hemispiralis* and *T. kiaochowensis* to stabilize the species names objectively, mainly because of the previous unavailability of type materials. By phylogenetic analysis and comparison with closely-related species, we infer that the ciliary tuft and details such as the commencement of the rightmost kinety in the lateral ciliary field are synapomorphies that may help clarify the systematics of *Tintinnopsis*-like taxa.

**Conclusion:** The redescriptions of three poorly known *Tintinnopsis* species, namely *T. hemispiralis*, *T. kiaochowensis*, and *T. uruguayensis* firstly revealed their ciliary patterns and rRNA sequences. This study expands knowledge and database of tintinnines and helps in identifying potential synapomorphies for future taxonomic rearrangements.

**Background**

Ciliated protists are among the most diverse and numerically important members of microzooplankton, and act as a trophic link in the microbial food web of aquatic ecosystems [1][2][3][4][5]. In particular, tintinnine ciliates are conspicuous due to the diversity of loricae produced by their cell propers. Tintinnines have been of great interest in the field of protistology because they (i) display distinct patterns of diversity and biogeography [6][7]; (ii) serve as bioindicators of water quality and hydrological circulation [8][9][10][11]; (iii) are prey for fish larvae and other small metazoans [12][13]; and (iv) can leave fossilized loricae that are useful in evolutionary studies [14][15].

There are approximately 1,000 extant tintinnine species classified almost entirely based on the shape and size of their loricae [16][17][18][19][20][21]. However, it is widely recognized that lorica features alone have shortcomings for determining taxonomic affiliations in this group of ciliates [22][23]. In some species, laboratory cultures have provided clear evidences that the lorica is polymorphic in response to
environmental factors or in different stages of the life cycle [24]. More recently, DNA sequencing of several closely-related species has revealed examples of polymorphic and cryptic species [25][26]. Thus, the current lorica-based taxonomy does not allow estimating tintinnine diversity accurately, and it does not provide a natural classification. Accordingly, several studies have incorporated more informative characters, namely, cytological and/ or molecular data, in tintinnine systematics (e.g., [27][28][29][30][31] [32][33][34][35][36]). Still, cell characters and DNA sequences are only known for about 3% and 10% of the described tintinnine morphospecies, respectively (e.g., [22][37]), and considerable efforts are needed to increase the availability of these types of information.

Arguably the most problematic taxon in tintinnine taxonomy is the genus *Tintinnopsis* Stein, 1867. This genus is known as artificial, given that it includes at least five distinct ciliary patterns [22][33][38] and more than ten clades that are non-monophyletic in rDNA sequence analyses [39]. Out of the about 140 *Tintinnopsis*-like morphospecies [19][20][21], about 60 have been recorded in China seas (e.g. [21][40][41] [42][43][44]), but only a few count with ciliature and/or sequence data [34][45][46][36][47][48]. Overall, *Tintinnopsis* will need subdivision once its type, *T. beroidea*, and other species are studied with modern methods [22][38][39].

The present study investigates the morphology and molecular phylogeny of three *Tintinnopsis* species, namely, *T. hemispiralis* Yin, 1956, *T. kiaochowensis* Yin, 1956, and *T. uruguayensis* Balech, 1948, which were collected from coastal waters of China. This work includes observations of specimens *in vivo* and after protargol staining as well as phylogenetic analyses of ribosomal RNA gene markers based on recommendations for tintinnine taxonomy [23] and common practices for other ciliates [49]. The aims of the present study are to combine lorica, cell proper, and molecular data in three *Tintinnopsis* species and to compare them with related taxa in order to find potential diagnostic features relevant in this problematic tintinnine taxon.

**Zoobank Registration**

The ZooBank registration number of the present work is: urn:lsid:zoobank.org:pub:38490F0B-183F-45AE-A053-80FCA6799716

**Results**

**Order Choreotrichida Small and Lynn, 1985**

**Suborder Tintinnina Kofoid and Campbell, 1929**

**Genus Tintinnopsis Stein, 1867**

*Tintinnopsis hemispiralis* Yin, 1956 (Figures 1A–E, 2A–J; Table 1)

**Terminology**
*Tintinnopsis hemispiralis* possesses a cluster of extremely long cilia that has only been reported for *Tintinnopsis subacuta* [50]. This character is here defined as follows.

**Ciliary tuft.** An extraordinary long tuft of cilia originated from densely arranged kinetids in the middle portion of the ventral kinety.

**Improved diagnosis (based on the type and neotype populations)**

Lorica 88–182 μm long, comprising a cylindrical, spiraled collar and an obconical bowl. Opening 34–59 μm in diameter. Cell proper elongate, obconical when fully extended, size *in vivo* 80–125 × 30–55 μm. Seven to 11 moniliform macronuclear nodules. On average 21 collar membranelles, of which four or five elongate into buccal cavity; one buccal membranelle. Ventral kinety composed of about 53 monokinetids, commences anteriorly to the second kinety of right ciliary field. Ciliary tuft about 150–250 μm long. Right and left ciliary fields consist of about ten kineties each. Lateral ciliary field comprises on average 15 kineties. Dorsal kinety composed of about 47 dikinetids. Posterior kinety with about 17 dikinetids, positioned below left ciliary field.

**Deposition of neotype and other voucher materials**

A protargol slide including the neotype (Figures 2F, G) was deposited in the Laboratory of Protozoology, Institute of Evolution and Marine Biodiversity, Ocean University of China (registration number: BY201805280101). One additional protargol slide was deposited in the same collection (registration number: BY201805280102).

**Redescription based on the Ningde population**

Lorica 143–182 μm long, comprises a cylindrical, truncated collar and an obconical bowl (Figures 1A, 2A–D). Opening 45–59 μm across. Ratio of lorica length to opening diameter 2.9–3.2:1. Collar 64–97 μm long, with three to five inconspicuous spiraled striations (Figures 1A, 2B). Bowl often slightly wider than opening (49–66 μm in diameter), about 68–88 μm long, with a posterior angle of 45° (Figures 1A, 2B, C, D). Wall of lorica heterogeneously agglutinated with mineral particles: collar slightly less agglutinated than bowl because adhered particles sparser and thinner (Figures 1A, 2B, C, D).

Cell proper elongate, obconical when fully extended, size 80–125 × 30–55 μm *in vivo*, and 65–119 × 31–58 μm after protargol staining (Figures 1B, 2E). Posterior portion of cell proper narrows gradually forming a peduncle with a branched posterior end, which is about 60–110 μm long and attaches to bottom of lorica (Figures 1B, 2D, E). Seven to 11 moniliform macronuclear nodules, each about 5–10 long and 5–9 μm wide; anterior nodule 17–21 μm posterior to the anterior cell end in protargol-stained specimens (Figures 1C, D, 2F, I, J). Micronuclei, striae, tentaculoids, accessory combs, a contractile vacuole, a cytopyge, and capsules not recognized. Cytoplasm colourless, with food vacuoles of various sizes containing ingested ovoidal microalgae (Figures 1B, 2E). Locomotion by irregular swim with rotation about main cell axis.
Somatic ciliary pattern composed of a ventral, a dorsal, and a posterior kinety and right, left, and lateral ciliary fields (Figures 1C–E, 2F–I). Kinetids of each ciliary row ostensibly connected by argyrophilic fibers (Figure 2F, G). Ventral kinety commences between collar membranelles and second kinety of right ciliary field (about 4–8 μm below the anterior end of cell) and curves leftwards before extending, parallel to kinetics of lateral ciliary field posteriorly; 39–66 μm long, with 41–61 monokinetids, consisting of three portions: (1) anterior portion comprised of eight to 14 kinetids about 0.5–1 μm apart; (2) middle portion consisting of 16–24 more densely arranged kinetids (with no measurable gap) with long cilia and forming the ciliary tuft, about 150–250 μm long in vivo; (3) posterior portion containing sparsely arranged monokinetids (more than 1 μm apart), extending posteriorly and terminating at about two thirds to three fourths of cell (Figures 1A–C, E, 2C, F, G). Right ciliary field comprises 9–11 kinetics, neighboring kinetics about 2–5 μm apart, each composed of about 5–18 widely spaced monokinetids and one anterior dikinetid; kinetids of first kinety more densely arranged; all kinetics commence at the same level (about 9 μm below the anterior end of cell), except for the first kinety which commences about 2 μm below the level of remaining kinetics (Figures 1C, E, 2F). Left ciliary field comprises 9–12 kinetics, neighboring kinetics about 2–5 μm apart; each kinety comprises about 9 μm below the anterior end of cell and is composed of one anterior dikinetid and 2–13 widely spaced monokinetids; the leftmost two or three kinetics always shorter, only including three to five kinetics (Figures 1C–E, 2G–I). Each basal body in left and right ciliary fields bears a cilium, the anterior one in each dikinetid being about 20 μm long in vivo and 10 μm long after protargol staining; the other cilia (on posterior kinety of dikinetids and on monokinetids) are about 3 μm long after protargol staining (Figures 1A, B, 2E, I). Lateral ciliary field consists of 11–20, relatively densely arranged monokinetidal kinetics, each of which commences about 9 μm below the anterior end of cell; kinetics in middle region always shorter than those at both ends of field (i.e., including only half the number of kinetics); cilia about 3 μm long after protargol staining (Figures 1D, E, 2G, H). Dorsal kinety commences about 5 μm below the anterior end of cell, about 5 and 10 μm apart from right and left ciliary field, respectively; curves towards the left-posterior portion; 66–97 μm long and composed of 35–56 dikinetids, with only the posterior basal body bearing a cilium about 8–10 μm long after protargol staining (Figures 1D, E, 2G, I). Posterior kinety usually commences below right portion of left ciliary field (29–41 μm below the anterior end of cell) and curves rightwards, terminating near posterior end of cell proper; 37–58 μm long and with 11–22 dikinetids; only the posterior basal body bearing a cilium about 8–10 μm long after protargol staining (Figures 1D, E, 2G–I).

Oral apparatus occupies anterior portion of cell. Adoral zone of membranelles closed; composed of 20–22 collar membranelles, four or five of which extend into buccal cavity; bases about 30 μm of longest membranelles; polykinetid structures could not be recognized; cilia of membranelles about 25–35 μm long (Figures 1A–E, 2A–E, J). Single buccal membranelle within buccal cavity, with polykinetid about 40 μm long (Figure 1C, E). Argyrophilic fibers originate in the proximal portions of the elongated collar membranelles and the buccal membranelle, and extend posteriorly; three thick fibers commencing from the middle of cell below right ciliary field and extending towards anterior part of cell; ends not observed due to insufficient staining (Figure 2F). Endoral membrane composed of a single row of basal bodies, extends in a semicircle across the peristomial field and right wall of buccal cavity (Figure 1C). An early
divider was observed with the oral primordium posterior to the ventral kinety and lateral ciliary field (Figure 2H).

*Tintinnopsis kiaochowensis* Yin, 1956 (Figures 3A–E, 4A–K; Table 1)

**Improved diagnosis (based on the type and neotype populations)**

Lorica 79–112 μm in length, composed of an irregular collar and an ellipsoidal bowl with a rounded posterior end, both separated by a constriction. Opening 30–71 μm in diameter. Cell proper obconical when fully extended, size *in vivo* about 60–95 × 35–50 μm. Two ellipsoidal macronuclear nodules. On average 16 collar membranelles, three of which extend into buccal cavity; one buccal membranelle. Ventral kinety with an average of 49 densely arranged monokinetids. Right, left, and lateral ciliary fields include, on average, 11, 10, and 16 kineties, respectively. Dorsal kinety composed of about 31 dikinetids. Posterior kinety composed of about 15 dikinetids, positioned below lateral ciliary field.

**Deposition of neotype and other voucher materials**

A protargol slide including the neotype (Figure 4H, I) was deposited in the Laboratory of Protozoology, Institute of Evolution and Marine Biodiversity, Ocean University of China (registration number: BY201805280201). One additional protargol slide was deposited in the same collection (registration number: BY201805280202).

**Redescription based on the Ningde population**

Lorica 79–112 μm in length, composed of an irregular collar and an ellipsoidal bowl (Figures 3A, 4A–C). Opening 44–71 μm in diameter; rim irregular. Ratio of lorica length to opening diameter 1.4–2:1. Collar 32–46 μm high, not flaring at the opening margin, occasionally slightly layered because of agglutinated particles arranged in horizontal rows (Figures 3A, 4A–C). Region between collar and bowl constricted, about 38–63 μm in diameter (Figures 3A, 4A, B). Bowl about 43–70 μm long and 57–83 μm across. Posterior end usually rounded to bluntly tapered (Figures 3A, 4A, B).

Cell proper elongate, obconical when fully extended, about 60–95 × 35–50 μm *in vivo*, and 46–65 × 38–64 μm after protargol stained. Posterior cell portion narrows successively forming a peduncle about 25 μm long and attached to the bottom of lorica (Figures 3A, D, 4B). Two ellipsoidal macronuclear nodules, each about 15–22 × 12–17 μm in protargol-stained specimens; anterior node 11–24 μm from the anterior cell end (Figure 3C). Micronuclei, striae, tentaculoids, accessory combs, a contractile vacuole, a cytopyge, and capsules not recognized. Cytoplasm colourless, usually with food vacuoles containing diatoms and ovoidal microalgae (Figures 3D, 4B). Locomotion by rotation about main cell axis.

Somatic ciliary pattern complex, with a ventral, a dorsal, and a posterior kinety, and right, left, and lateral ciliary fields (Figures 3B, C, E, 4H–K). Ventral kinety 22–37 μm long, commences between collar membranelles and third or fourth kinety of right ciliary field (about 4 μm below the anterior end of cell), anterior third curves leftwards before extending parallel to kineties of lateral ciliary field posteriorly; 43–
56 densely arranged monokinetids (Figures 3B, E, 4H). Right ciliary field comprising 10–13 kineties about 2–5 μm apart, the space between the leftmost five to six kineties wider than others; all kineties commence 6–13 μm below the anterior end of cell; composed of 5–14 widely spaced monokinetids and one anterior dikinetid, except for: (i) the first kinety almost parallel to ventral kinety, with two or three anterior dikinetids and eight to 12 monokinetids, more densely arranged than other kineties in right ciliary field; and (ii) the second kinety parallel to rest of kineties, with an angle of about 20° with the first kinety, including four or five monokinetids and two anterior dikinetids (Figures 3B, E, 4H). Left ciliary field comprises 9–11 kineties about 2–5 μm apart, each kinety commencing 6–13 μm below the anterior end of cell and comprised of two to eight widely spaced monokinetids and one anterior dikinetid; the number of kinetids of leftmost kinety always minimum (i.e., three or four). Each basal body in left and right ciliary fields bears a cilium, with the cilium on anterior basal body of each dikinetid being about 15 μm long in vivo and 5 μm long after protargol staining; other cilia (on posterior basal body of each dikinetid and on monokinetids) are about 1 μm long after protargol staining (Figures 3A, D, 4C, H–J). Lateral ciliary field with 13–19 monokinetidal kineties of similar length, each of which commences 6–13 μm below the anterior end of cell, except for the rightmost kinety that commences anteriorly to the second or third kinety of right ciliary field, about 4 μm below the anterior cell end, with the anterior portion curving rightwards before extending towards posterior part; cilia about 2 μm long after protargol staining (Figures 3B, C, E, 4H, I). Dorsal kinety 29–53 μm long, comprises 25–37 dikinetids, commences about 4 μm below the anterior end of cell, about 5 μm from right ciliary field and 13 μm from left ciliary field apart, and extends to the left before curving towards posterior end of cell (Figures 3C, E, 4J, K). Posterior kinety 21–29 μm long, consists of 11–18 dikinetids, commences below lateral ciliary field (28–49 μm below the anterior end of cell) and extends to posterior end of cell (Figures 3C, E, 4J). Cilia of dorsal and posterior kinety are insufficiently stained.

Oral apparatus occupies anterior portion of cell. Adoral zone of membranelles closed, composed of 16–18 collar membranelles with cilia about 25–35 μm long, three of which extend into buccal cavity; bases about 30 μm of longest membranelles; kinetal structures of membranelles could not be recognized (Figures 3A–E, 4E–G, H). Single buccal membranelle in buccal cavity, with polykinetid about 20 μm long (Figures 3B, E, 4E). Endoral membrane composed of a single row of basal bodies, extends in a semicircle across the peristomial field and right wall of buccal cavity (Figures 3B, C, 4E, F). Argyrophilic fibers associated with oral apparatus insufficiently impregnated to be observed.

*Tintinnopsis uruguayensis* Balech, 1948 (Figures 5A–D, 6A–J; Table 1)

**Improved diagnosis (based on the type and present populations)**

Lorica 50–73 μm long, bullet-like with a flared collar and a posterior process about 8–10 μm long. Opening 22–42 μm in diameter, with an irregular rim. Cell proper obconical when fully extended, size in vivo about 25–50 μm × 20–30 μm. Two macronuclear nodules. On average 18 collar membranelles, of which three or four extend into buccal cavity; one buccal membranelle. Ventral kinety composed of about 20 monokinetids. Right and left ciliary fields consist of about seven kineties each. Lateral ciliary field
comprises on average 12 kineties. Dorsal kinety with about 21 dikinetids. Posterior kinety with about eight dikinetids, posterior to lateral ciliary field.

**Deposition of voucher materials**

Two protargol slides with voucher specimens were deposited in the Laboratory of Protozoology, Institute of Evolution and Marine Biodiversity, Ocean University of China (registration numbers: BY201811120101 and BY201811120102).

**Redescription based on the Qingdao population**

Lorica 50–73 μm long, composed of a flared collar about 15 μm long with a jagged rim, and an ovoidal bowl about 32–52 μm long and 25–41 μm wide (Figures 5A, 6A–C). Opening diameter 24–42 μm. Region between collar and bowl narrowed, about 17–29 μm in diameter (Figures 5A, 6A–C). Posterior end projected, about 10 μm long (Figures 5A, 6A). Lorica wall densely agglutinated with mineral particles (Figures 5A, 6A–C).

Cell proper obconical when fully extended, about 25–50 μm × 20–35 μm in size *in vivo*, and 25–56 × 18–28 μm in size after protargol staining. Posterior end of cell proper becomes spherical when escaped from lorica (Figure 6D). Two ellipsoidal (occasionally elongated) macronuclear nodules, 6–14 × 4–10 μm in size after protargol staining; anterior nodule 4–9 μm posteriorly to the anterior cell end after protargol staining (Figures 5A–C, 6E, H). Micronuclei, striae, tentaculoids, accessory combs, a contractile vacuole, a cytopyge, and capsules not recognized. Cytoplasm colourless, with food vacuoles of various sizes containing ingested microalgae (Figure 6D). Locomotion by rotation about main cell axis. Disturbed individuals retract into lorica with motionless membranelles bent to centre of peristomial field.

Somatic ciliary pattern composed of a ventral, a dorsal, and a posterior kinety, and right, left, and lateral ciliary fields (Figures 5B–D, 6E–I). Ventral kinety 14–35 μm long with 17–28 monokinetids, commences between collar membranelles and first kinety of right ciliary field, about 2 μm below the anterior end of cell, curves leftwards before extending parallel to kinetics of lateral ciliary field posteriorly (Figures 5B, D, 6E). Right ciliary field comprises 7–8 kinetics, 1–3 μm apart; all kinetics commence at the same level (about 2 μm below the anterior end of cell), except for the first kinety that commences about 1 μm below the level of remaining kinetics; the second kinety always shorter, i.e. only comprising two or three kinetids; others composed of 6–7 widely spaced monokinetids and one anterior dikinetid, except first kinety comprised of two to four monokinetids and two or three anterior dikinetids; first kinety usually commences below anterior portion of ventral kinety (Figures 5B, D, 6E, I). Left ciliary field comprises 6–8 kinetics; each kinety commences about 2 μm below the anterior end of cell and is composed of one anterior dikinetid and 1–7 widely spaced monokinetids, with decreasing length from right to left (Figures 5C, D, 6F, G). Each basal body in left and right ciliary fields bears a cilium, with the cilium on anterior basal body of each dikinetid being about 5 μm long *in vivo* and after protargol staining; other cilia (on posterior basal body of each dikinetid and on monokinetids) are about 1 μm long after protargol staining (Figures 5A, 6E–J). Lateral ciliary field commences about 2 μm below the anterior end of cell, with 9–16
monokinetidal kineties of similar length, with cilia about 1 μm long after protargol staining (Figures 5B, D, 6E). Dorsal kinety commences about 2 μm below anterior cell end, about 2 and 3 μm apart from right and left ciliary fields, respectively, curves towards the left-posterior portion of cell; 21–41 μm long, consists of 17–29 dikinetids, with only the posterior basal body bearing a cilium about 3–5 μm long after protargol staining (Figures 5C, D, 6F, G). Posterior kinety usually commences below the middle kinety of the left ciliary field (12–21 μm below the anterior end of cell) and curves leftwards; 11–22 μm long, consists of 7–9 dikinetids, with only the posterior basal body bearing a cilium about 3–5 μm long after protargol staining (Figures 5C, D, 6F).

Oral apparatus occupies anterior portion of cell. Adoral zone of membranelles closed; composed of 18 or 19 collar membranelles with cilia up to about 20–25 μm long, three or four of which extend into buccal cavity; bases about 10 μm of longest membranelles; polykinetid structures could not be recognized (Figures 5A–D, 6A, D, H). Single buccal membranelle in buccal cavity, with polykinetid about 8 μm long (Figures 5B, D, 6H). Argyrophilic bers insufficiently impregnated to be observed. Endoral membrane not recognized. One middle divider was observed with the oral primordium located left of ventral kinety and posterior to the lateral ciliary field (Figure 6J).

Neotypification

The neotypes of *Tintinnopsis hemispiralis* and *T. kiaochowensis* are designated because (i) no type materials are known to be deposited; (ii) the original descriptions are restricted to lorica features, while the present redescriptions include also cytological and molecular analyses; and (iii) the type locality of the original populations is Qingdao, East China, with no further details [44]. The type location of the two species is nearby the collection site of the present populations (Meng Bay, Ningde, East China; detailed information provided in ‘Materials and Methods’), thus meeting the requirement of Article 75.3.6 of the International Code of Zoological Nomenclature [51]. Protargol slides containing the neotype specimens were deposited (see ‘Deposition of neotype and other vouched materials’), thus meeting the requirements of Article 75.3.7 of the Code [51]. A neotype is not established for *T. uruguayensis* because the type location corresponds to a different ocean basin [52].

Sequence comparison and phylogenetic analyses

For the three species investigated, the length, G+C content and GenBank accession numbers of the SSU rDNA, ITS1-5.8S rDNA-ITS2 and LSU rDNA sequences are compiled in Table 2. For each of the three loci and concatenated sequences, the topologies of the Maximum Likelihood (ML) and Bayesian Inference (BI) trees were similar and therefore only the ML trees are shown (Figures 7, 8, 9, S1). *Tintinnopsis hemispiralis* forms a fully-supported clade with *T. subacuta* (EU399541; [53]) based on SSU rDNA; both sequences are 99.3% similar. Based on ITS1-5.8S-ITS2, a sequence previously obtained for this species in Qingdao, China (KU715813; [48]) groups with our sequence, and both are 96.2% similar. *Tintinnopsis kiaochowensis* forms a fully-supported clade with *T. everta* (MG461220; [33]) based on SSU rDNA, and both sequences are 99.0% similar. The newly sequenced population of *T. uruguayensis* forms a fully-supported clade with the North-Atlantic population of the same species, based on both SSU rDNA and
LSU rDNA (JN831838 and JN831923; [25]); the two populations are 100% identical in both markers. The concatenated tree (Figure S1, Table S1) shows similar relationships than SSU rDNA, except that Tintinnina were inferred as non-monophyletic. This inference is probably artifactual given the well-known monophyly of this suborder [22][39].

Discussion

*Tintinnopsis hemispiralis*

Comparison with other populations

The specimens studied here match *Tintinnopsis hemispiralis* in lorica size and shape [44]. The lorica dimensions reported in the original description (length = 88–164 μm, opening diameter = 34–53 μm; [44]) overlap with those of our specimens (length = 143–182 μm, opening diameter = 45–59 μm; Table 1). The originally described population and our specimens also match in a lorica composed of a cylindrical, spiraled collar and an obconical bowl (Figures 1A, 2B–D). One ITS1-5.8S rDNA-ITS2 sequence labeled as *T. hemispiralis* in GenBank [48] presents a relatively high divergence (3.8%) compared to our sequence; conspecificity of both populations cannot be confirmed.

Comparison with similar species

Four congeners, namely *Tintinnopsis cochleata* (Brandt, 1906) Laackmann, 1913, *Tintinnopsis directa* Hada, 1932, *Tintinnopsis gracilis* Kofoid and Campbell, 1929, and *Tintinnopsis tubulosoides* Meunier, 1910, are similar to our specimens in an elongated lorica with a spiraled collar. *Tintinnopsis cochleata* differs from our specimens in a sub-hemispherical posterior end and 13 (vs. 3–5) spiral striations in the collar of the lorica [16]. *Tintinnopsis directa* can be separated from our population by a swollen, ovoid bowl and rounded posterior end of the lorica (vs. coniform, pointed; [54]). *Tintinnopsis gracilis* differs from our specimens by smaller lorica size (110–135 μm long vs. 143-182 long) and spiraled striation absent in collar portion (vs. 3–5 obvious spiraled striations, see Figure 2B) [19]. *Tintinnopsis tubulosoides* differs from our specimens in a smaller lorica size (91 μm long and 33 μm in opening diameter, based on the illustration included in the original description) and two (vs. 7–11) macronuclear nodules [55]. For these species, rDNA sequences have been reported only for *T. tubulosoides* (AF399111–AF399020; [56]), which shows a distant relationship to *T. hemispiralis* (Figures 7, 8).

Regarding cell features, *T. hemispiralis* resembles *Tintinnopsis subacuta* Jörgensen, 1899 in having a ventral kinety associated with the extraordinarily long ciliary tuft that extends outside of the lorica [50]. Both species are also similar in having multiple moniliform macronuclear nodules [50], which differ from the common finding of only two macronuclear nodules in other *Tintinnopsis* species (e.g., [22][27][34]). The two species cluster together based on SSU rDNA (Figure 7), which suggests that the ciliary tuft and multiple moniliform macronuclear nodules are synapomorphies of this clade and may be important for a future reclassification of *Tintinnopsis* species. Despite the close relationship between *T. hemispiralis* and *T. subacuta*, the latter can be distinguished from our specimens by a lorica with a swollen, ovoid (vs.
obconical) bowl in the original description [57] and the micrograph of the sequenced specimen [53]. The SSU rDNA divergence for both species, although small (0.7%), is consistent with interspecific variation in this conserved marker [25].

*Tintinnopsis kiaochowensis*

Comparison with type population

The specimens studied here match *Tintinnopsis kiaochowensis* in lorica size and shape [44]. The lorica dimensions reported in the original description (length = 95–108 μm, opening diameter = 30–52 μm; [44]) overlap with those of our specimens (length = 79–112 μm, opening diameter = 44–71 μm; Table 1). Our specimens also resemble to those originally described in a lorica with a cylindrical collar and an ellipsoidal bowl with a constricted connection. However, our specimens differ from the original population in the rounded posterior end of the lorica (vs. obconical) and in the agglutinated particles forming horizontal rows on the collar (vs. both on collar and bowl) [44].

Comparison with similar species

*Tintinnopsis kiaochowensis* differs from other *Tintinnopsis* species by its peculiar lorica shape, i.e. swollen bowl divided from a non-flaring collar by a constriction. Compared to our specimens, the most similar species is *Tintinnopsis compressa* Daday, 1887. However, *T. compressa* can be separated from our specimens by having a smaller lorica size (45 vs. 79–112 μm in length; 26 vs. 44–71 μm in opening diameter), a flared lorica collar (vs. not flared), and a less obvious constriction between the lorica collar and bowl [18].

*Tintinnopsis kiaochowensis* is similar to *Tintinnopsis everta* Kofoid and Campbell, 1929 based on SSU rDNA (Figure 7) and cytological characters [33], including: (i) elongated anterior portion of the ventral kinety, which forms a curvature above the third, occasionally the fourth, kinety of the right ciliary filed; (ii) elongated anterior portion of the rightmost kinety of lateral ciliary field, which forms a curvature above the second or third kinety of the right ciliary field (with the ventral kinety in between); and (iii) first four to six kineties of the right ciliary field very widely spaced. However, unique cytological features observed in *T. everta* (the large distance between the collar membranelles and the somatic ciliary fields as well as the position of the posterior kinety; [33]) are not present in *T. kiaochowensis*. Both species also show a different lorica morphology (campanulate lorica with a funnel-shaped collar vs. ellipsoidal bowl and non-flaring collar, respectively) and interspecies-level divergence in SSU rDNA (1%; [25]).

*Tintinnopsis uruguayensis*

Comparison with other populations

This species was first described by Balech [52] based on the lorica features of specimens collected in the Southwest Atlantic Ocean. The lorica dimensions reported in the original description (length = 54–63 μm, opening diameter = 22–27 μm; [52]) overlap with those of our specimens (length = 50–73 μm, opening
diameter = 24–42 μm; Table 1), and both populations match in the characteristic bullet-like shape with a flared collar and a posterior process. Our population presents no divergence in SSU rDNA and LSU rDNA when compared against Long Island Sound specimens of similar lorica features [25].

Comparison with similar species

In terms of a small, bullet-like lorica, three congeners, namely *Tintinnopsis baltica* Brandt, 1896, *Tintinnopsis fimbriata* Meunier, 1919, and *Tintinnopsis meunieri* Kofoid and Campbell, 1929, can be compared to our population. *Tintinnopsis baltica* has a similar lorica shape, but can be separated from *T. uruguayensis* by the absence (vs. presence) of a protruding posterior end [58]. Laval-Peuto & Brownlee [59] provided a diagram of the ciliary pattern of *T. baltica*, which is similar to our specimens in the number of kineties in the right, left, and lateral ciliary fields and the presence of only 2–3 kinetids in the second kinety of right ciliary field, but differs in a shorter ventral kinety. The distant phylogenetic relationship between *T. uruguayensis* and *T. baltica* based on SSU rDNA and LSU rDNA (Figures 7, 9) also separates both species. *Tintinnopsis fimbriata* differs from *T. uruguayensis* by a shorter collar (10 μm vs. up to 20 μm) and a wider bowl (40–50 μm vs. 25–41 μm) [60]. Based on cytological data [27], *T. fimbriata* also differs from the latter in having less kineties in the left ciliary field (4–6 vs. up to 9) and lateral ciliary field (11–14 vs. up to 17). The SSU rDNA sequence labeled as *T. fimbriata* in GenBank (Figure 7) has been considered a misidentification [39] and is thus not considered in this comparison. *Tintinnopsis meunieri* differs from *T. uruguayensis* in a larger opening diameter (60 μm vs. 24–42 μm) [19].

Conclusion

*Tintinnopsis hemispiralis*, *T. kiaochowensis* and *T. uruguayensis* show hard, fully agglomerated loricae and the most complex pattern of somatic ciliature known for the genus, i.e. a right, left and lateral ciliary field as well as a ventral, dorsal and posterior kinety [22]. However, the three species show differences in the lorica outline and the number, structure and arrangement of somatic kineties (Figures 1–6; Table 1), and species-level divergence in rRNA genes [25][26]. Their distant position and intertwining with other genera in phylogenetic trees (Figures 7, 8, 9) confirm, once again, the non-monophyly of the genus *Tintinnopsis* [22][38][39].

*Tintinnopsis* cannot be revised at present, as its type species and most other tintinnine species have not been studied cytologically or genetically [23]. Our work is important to increase the number of tintinnine species investigated with modern methods, which also helps in identifying potential synapomorphies for future taxonomic rearrangements. Our data show the potential taxonomic relevance of (i) details of the somatic ciliary pattern, including the anterior parts of the ventral kinety and the rightmost kinety of the lateral ciliary field [33]; and (ii) the presence of a ciliary tuft and multiple moniliform macronuclear nodules. Our paper contributes important information on the non-monophyletic *Tintinnopsis* and it thus helps to fill the gaps in modern tintinnine taxonomy.

Methods
Sample collection and morphological analysis

*Tintinnopsis hemispiralis* and *Tintinnopsis kiaochowensis* were collected from surface coastal waters in Meng Bay, Ningde, Fujian Province, China (25°54'24''N 119°40'22''E; temperature = 25°C; salinity = 30) on May 28, 2018 (Figure 10A, B); *Tintinnopsis uruguayensis* was collected from surface coastal waters off Qingdao, Shandong Province, China (36°03'35''N 120°18'53''E; temperature = 22°C; salinity = 30) on November 12, 2018 (Figure 10A, C).

The cells were isolated by using micropipettes under a stereo microscope (Olympus SZX2-TR30, Tokyo, Japan) at 45× magnification. Live cells were studied with bright-field and differential interference contrast microscopy at 100–1000× total magnification. Loricae were measured from living cells at magnifications 100–400×. The loricae were separated from their cell proper with an eyebrow brush after fixation in Bouin’s solution. Protargol staining followed Wilbert [61]. The protargol powder was manually synthesized following Pan et al. [62]. Protargol-stained specimens were measured at 1,000× magnification. Drawings were made at a 1,000× magnification with the aid of a camera lucida. Identities were based on original descriptions [44][52] and other tintinnine bibliography mentioned above. Terminology and classification follow Agatha & Riedel-Lorjé [63] and Adl et al. [64], respectively.

DNA extraction, PCR amplification and sequencing

Because most tintinnine species are not amenable to culture, clonal cultures could not be established. Thus, we applied common criteria to verify that field-isolated specimens were not confounded with other species (e.g. as done by Gruber et al. [33]): the three species were distinguished by careful evaluation of their morphological features and loria size in vivo, and the absence of potentially confounding, co-occurring species was confirmed with further analyses of loricae and protargol-stained cells. For each species, a single specimen was isolated at 400× magnification and washed five times with 0.22-μm filtered sample water. DNA extraction, PCR amplification and sequencing were done as detailed by Bai et al. [31], except for some of the primers utilized. PCR amplification of the SSU rDNA was performed with the primers 82F (5'-GAA ACT GCG AAT GGC TC-3'; [65]) and either 5.8s-R (5'-CTG ATA TGC TTA AGT TCA GCG G-3'; [66]) for *Tintinnopsis uruguayensis* or 18s-R (5'-TGA TCC TTC TGC AGG TTC ACC TAC-3'; [67]) for the other two species. A fragment containing the ITS1, 5.8S rDNA and ITS2 regions was amplified with the primers 5.8s-F (5'-GTA GGT GAA CCT GCG GAA GGA TC-3') and 5.8s-R (5'-CTG ATA TGC TTA AGT TCA GCG G-3') [66].

Sequences were assembled and analysed as reported before [31]. In brief, phylogenetic analyses were done separately for SSU rDNA, ITS1-5.8S rDNA-ITS2 and LSU rDNA, as well as after concatenating the three sequence markers. The analyses incorporated additional ciliate sequences were obtained from GenBank and used *Halteria grandinella* and hypotrichs as outgroup taxa. Sequences were aligned with Muscle 3.7 [68]. Maximum likelihood analyses were done with RAxML v. 8 [69], using the GTRGAMMA model and 1,000 bootstraps. Bayesian Inference analyses were done with MrBayes v.3.2.6 [70], using the GTR + I + Γ model, 6,000,000 generations with a sample frequency of 100 generations and a burn-in of 6,000 trees. Estimates of sequence similarity were done in MEGA 7.0 [71].
List Of Abbreviations

BM: buccal membranelle; CM: collar membranelle; CV: coefficient of variation in %; DK: dorsal kinety; EM: endoral membrane; LA: lateral ciliary field; LF: left ciliary field; M: median; Ma: macronuclear nodules; Max: maximum; Mean: arithmetic mean; Min: minimum; N: number of specimens examined; PCM: prolonged collar membranelles; PK: posterior kinety; RF: right ciliary field; SD: standard deviation; VK: ventral kinety.

Declarations

Ethics approval and consent to participate

No field permissions were necessary to collect the samples for this study. The authors declared that the experimental research on the protists described in this paper was in compliance with institutional, national and international guidelines.

Consent for publication

Not applicable.

Availability of data and materials

Sequence data is available in GenBank (Accession Numbers: MT435060–MT435062, MT435073–MT435078). Permanent slides containing the protargol-stained specimens of *Tintinnopsis hemispiralis*, *T. kiaochowensis*, and *T. uruguayensis* with registration numbers of BY201805280101, BY201805280102, BY201805280201, BY201805280202, BY201811120101, and BY201811120102 are Laboratory of Protozoology, Institute of Evolution and Marine Biodiversity, Ocean University of China.

Competing interests

The authors declare that they have no competing interests.

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Authors’ contribution

XH conceived and guided the study. YB, RW, and WS conducted sampling and performed laboratory work. XH, WS, LL, and YB identified the species. YB and LS did and interpreted the sequence similarity and phylogenetic analyses. YB drafted the manuscript, and WS, LL, LS and XH made further revisions. All authors read and approved this manuscript.
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Tables

Table 1. Morphometric data of *Tintinnopsis hemispiralis*, *T. kiaochowensis*, and *T. uruguayensis* (measurements in μm). Lorica data are based on live specimens, and other data are based on protargol-stained specimens. Abbreviations:
| Characters                        | Species name      | Min  | Max  | Mean | M   | SD  | CV  | N  |
|----------------------------------|-------------------|------|------|------|-----|-----|-----|----|
| Lorica, total length             | T. *hemispiralis* | 143  | 192  | 161.1| 161 | 12.9| 8.0 | 15 |
|                                  | T. *kiaochowensis*| 79   | 112  | 89.9 | 90  | 8.2 | 9.1 | 12 |
| Lorica, total length             | T. *uruguayensis* | 50   | 73   | 62.3 | 62  | 7.7 | 12.4| 15 |
| Lorica, bowl width               | T. *hemispiralis* | 49   | 66   | 55.9 | 57  | 5.1 | 9.2 | 15 |
|                                  | T. *kiaochowensis*| 57   | 81   | 66.7 | 65  | 7.4 | 11.0| 12 |
| Lorica, bowl width               | T. *uruguayensis* | 25   | 41   | 32.6 | 32  | 4.5 | 13.7| 15 |
| Lorica, bowl length              | T. *hemispiralis* | 68   | 88   | 77.8 | 76  | 6.6 | 8.5 | 15 |
|                                  | T. *kiaochowensis*| 43   | 70   | 51.3 | 49  | 7.2 | 14.1| 12 |
| Lorica, collar length            | T. *uruguayensis* | 32   | 63   | 48.8 | 47  | 7.7 | 15.9| 12 |
| Lorica, opening diameter         | T. *hemispiralis* | 45   | 59   | 52.6 | 54  | 4.5 | 8.6 | 15 |
|                                  | T. *kiaochowensis*| 44   | 71   | 55.1 | 53  | 7.5 | 13.6| 12 |
| Lorica, length: opening diameter | T. *uruguayensis* | 24   | 42   | 33.1 | 33  | 5.4 | 16.4| 15 |
| ratio                            | T. *hemispiralis* | 2.9  | 3.2  | 3.1  | 3.1 | 0.1 | 2.4 | 15 |
|                                  | T. *kiaochowensis*| 1.4  | 2.1  | 1.6  | 1.6 | 0.2 | 10.5| 12 |
| Lorica, narrowed portion diameter| T. *uruguayensis* | 1.6  | 2.3  | 1.9  | 1.9 | 0.2 | 8.2 | 15 |
|                                  | T. *kiaochowensis*| 38   | 63   | 48.8 | 47  | 7.7 | 15.9| 12 |
| Lorica, total length: narrowed   | T. *hemispiralis* | 1.4  | 2.2  | 1.9  | 2.0 | 0.3 | 16.2| 12 |
| portion diameter                 | T. *uruguayensis* | 1.8  | 3.2  | 2.6  | 2.6 | 0.4 | 16.0| 15 |
| Cell proper length               | T. *hemispiralis* | 65   | 119  | 95.1 | 95  | 15.0| 15.8| 15 |
|                                  | T. *kiaochowensis*| 46   | 65   | 55.9 | 58  | 5.8 | 10.4| 12 |
| Cell proper width                | T. *uruguayensis* | 25   | 56   | 31.5 | 30  | 7.4 | 23.5| 15 |
|                                  | T. *hemispiralis* | 31   | 58   | 44.5 | 47  | 8.3 | 18.6| 15 |
|                                  | T. *kiaochowensis*| 38   | 64   | 45.5 | 64  | 6.5 | 14.4| 12 |
| Macronuclear nodules, number      | T. *uruguayensis* | 18   | 28   | 22.5 | 22  | 3.0 | 13.3| 15 |
|                                  | T. *hemispiralis* | 7    | 11   | 9.2  | 9   | 1.0 | 11.0| 15 |
|                                  | T. *kiaochowensis*| 2    | 2    | 2.0  | 2   | 0.0 | 0.0 | 12 |
|                                  | T. *uruguayensis* | 2    | 2    | 2.0  | 2   | 0.0 | 0.0 | 15 |
| Metric                                      | Species            | Length 1 | Length 2 | Length 3 | Length 4 | Length 5 | Length 6 | Length 7 | Length 8 | Length 9 | Length 10 | Length 11 | Length 12 |
|--------------------------------------------|--------------------|----------|----------|----------|----------|----------|----------|----------|----------|----------|-----------|-----------|-----------|
| Macronuclear nodules, length               | *T. hemispiralis*  | 5        | 10       | 8.7      | 9        | 1.3      | 14.9     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 6                  | 14       | 8.3      | 7        | 2.3      | 27.2     | 15       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 15                 | 22       | 18.4     | 18       | 2.4      | 12.8     | 12       |          |          |          |           |           |           |
| Macronuclear nodules, width                | *T. hemispiralis*  | 5        | 9        | 7.4      | 7        | 0.9      | 12.3     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 12                 | 17       | 14.3     | 15       | 1.6      | 11.3     | 12       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 6                  | 14       | 8.3      | 7        | 2.3      | 27.2     | 15       |          |          |          |           |           |           |
| Anterior cell end to anterior macronucleus nodule, distance | *T. hemispiralis*  | 17       | 21       | 19.1     | 19       | 1.8      | 9.4      | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 11                 | 24       | 19.2     | 18       | 4.1      | 21.6     | 12       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 4                  | 9        | 6.3      | 6        | 1.4      | 22.1     | 15       |          |          |          |           |           |           |
| Ventral kinety, length                     | *T. hemispiralis*  | 39       | 66       | 56.6     | 55       | 5.2      | 9.3      | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 22                 | 37       | 32.3     | 34       | 4.7      | 14.5     | 12       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 14                 | 56       | 48.6     | 48       | 4.0      | 8.2      | 12       |          |          |          |           |           |           |
| Ventral kinety, number of kinetids         | *T. hemispiralis*  | 41       | 61       | 52.5     | 52       | 5.1      | 9.7      | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 43                 | 56       | 48.6     | 48       | 4.0      | 8.2      | 12       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 17                 | 28       | 20.3     | 21       | 2.3      | 11.3     | 15       |          |          |          |           |           |           |
| Ventral kinety, distance to anterior end of cell | *T. hemispiralis*  | 4        | 8        | 5.5      | 5        | 0.8      | 15.3     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 3                  | 5        | 4.1      | 4        | 0.5      | 12.6     | 12       |          |          |          |           |           |           |
| Dorsal kinety, length                      | *T. hemispiralis*  | 66       | 97       | 88.8     | 90       | 6.1      | 6.9      | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 29                 | 53       | 43.8     | 43       | 6.5      | 14.8     | 12       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 21                 | 41       | 26.5     | 24       | 5.5      | 20.8     | 15       |          |          |          |           |           |           |
| Dorsal kinety, number of kinetids          | *T. hemispiralis*  | 35       | 56       | 46.5     | 43       | 4.7      | 10.2     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 25                 | 37       | 31.2     | 31       | 4.4      | 14.3     | 12       |          |          |          |           |           |           |
| *T. uruguayensis*                          | 17                 | 29       | 21.1     | 20       | 3.5      | 16.7     | 15       |          |          |          |           |           |           |
| Dorsal kinety, distance to right ciliary field | *T. hemispiralis*  | 4        | 6        | 4.5      | 4        | 0.6      | 14.1     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 4                  | 6        | 4.7      | 5        | 0.7      | 14.0     | 12       |          |          |          |           |           |           |
| Dorsal kinety, distance to left ciliary field | *T. hemispiralis*  | 8        | 14       | 10.4     | 11       | 1.1      | 10.8     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 11                 | 21       | 13.2     | 13       | 2.6      | 19.7     | 12       |          |          |          |           |           |           |
| Dorsal kinety, distance to anterior end of cell | *T. hemispiralis*  | 2        | 5        | 3.2      | 3        | 0.7      | 21.8     | 15       |          |          |           |           |           |
| *T. kiaochowensis*                         | 4                  | 7        | 4.9      | 5        | 1.0      | 20.9     | 15       |          |          |          |           |           |           |
| Dorsal kinety, distance to anterior end of cell | *T. hemispiralis*  | 3        | 6        | 4.4      | 5        | 0.9      | 20.4     | 12       |          |          |           |           |           |
|                           | T. uruguayensis | 2   | 3   | 2.3 | 2  | 0.5 | 20.9 | 15 |
|---------------------------|-----------------|-----|-----|-----|----|-----|------|----|
|                           | T. hemispiralis | 37  | 58  | 45.9| 45 | 5.3 | 11.5 | 15 |
|                           | T. kiaochowensis| 21  | 29  | 25.5| 27 | 3.1 | 12.2 | 11 |
| **Posterior kinety, length** |                 |     |     |     |    |     |      |    |
|                           | T. uruguayensis | 11  | 22  | 13.7| 12 | 2.9 | 21.0 | 15 |
|                           | T. hemispiralis | 11  | 22  | 17.0| 17 | 2.7 | 15.7 | 15 |
|                           | T. kiaochowensis| 11  | 18  | 15.4| 15 | 2.0 | 12.8 | 11 |
| **Posterior kinety, number of kinetids** |                 |     |     |     |    |     |      |    |
|                           | T. uruguayensis | 11  | 22  | 17.0| 17 | 2.7 | 15.7 | 15 |
|                           | T. hemispiralis | 11  | 18  | 15.4| 15 | 2.0 | 12.8 | 11 |
|                           | T. kiaochowensis| 7   | 9   | 8.0 | 8  | 0.5 | 6.7  | 15 |
| **Posterior kinety, distance to anterior end of cell** | T. uruguayensis | 29  | 41  | 34.9| 34 | 3.4 | 9.7  | 15 |
|                           | T. hemispiralis | 28  | 49  | 36.0| 35 | 5.3 | 14.6 | 12 |
|                           | T. kiaochowensis| 29  | 41  | 34.9| 34 | 3.4 | 9.7  | 15 |
| **Right ciliary field, number of kineties** | T. uruguayensis | 11  | 18  | 15.4| 15 | 2.0 | 12.8 | 11 |
|                           | T. hemispiralis | 11  | 18  | 15.4| 15 | 2.0 | 12.8 | 11 |
|                           | T. kiaochowensis| 9   | 11  | 9.7 | 9  | 0.8 | 8.4  | 15 |
| **Longest kinety in right field, length** | T. uruguayensis | 16  | 26  | 19.7| 19 | 2.5 | 12.7 | 15 |
|                           | T. hemispiralis | 19  | 28  | 24.3| 25 | 2.5 | 10.4 | 12 |
|                           | T. kiaochowensis| 16  | 26  | 19.7| 19 | 2.5 | 12.7 | 15 |
| **Longest kinety in right field, number of kinetids** | T. uruguayensis | 15  | 19  | 16.6| 16 | 1.0 | 5.9  | 15 |
|                           | T. hemispiralis | 11  | 15  | 12.4| 12 | 1.1 | 8.7  | 12 |
|                           | T. kiaochowensis| 11  | 15  | 12.4| 12 | 1.1 | 8.7  | 12 |
| **Shortest kinety in right field, length** | T. uruguayensis | 7   | 8   | 7.3 | 7  | 0.5 | 6.3  | 15 |
|                           | T. hemispiralis | 6   | 16  | 10.7| 11 | 2.0 | 18.5 | 15 |
|                           | T. kiaochowensis| 7   | 8   | 7.3 | 7  | 0.5 | 6.3  | 15 |
| **Shortest in right field, number of kinetids** | T. uruguayensis | 6   | 11  | 8.6 | 8  | 2.2 | 18.5 | 15 |
|                           | T. hemispiralis | 6   | 7   | 6.5 | 7  | 0.5 | 8.0  | 12 |
|                           | T. kiaochowensis| 6   | 11  | 8.6 | 8  | 2.2 | 18.5 | 15 |
| **Left ciliary field, number of kineties** | T. uruguayensis | 9   | 12  | 9.8 | 9  | 1.0 | 10.3 | 15 |
|                           | T. hemispiralis | 9   | 11  | 9.9 | 10 | 0.7 | 6.7  | 12 |
|                           | T. kiaochowensis| 9   | 12  | 9.8 | 9  | 1.0 | 10.3 | 15 |
| **Longest kinety in left field, length** | T. uruguayensis | 6   | 8   | 6.6 | 6  | 0.8 | 12.5 | 15 |
|                           | T. hemispiralis | 17  | 28  | 23.3| 24 | 3.3 | 14.0 | 15 |
|                           | T. kiaochowensis| 11  | 16  | 12.5| 12 | 1.4 | 11.6 | 12 |
| **Longest kinety in left field, number of kinetids** | T. uruguayensis | 7   | 11  | 9.1 | 9  | 1.1 | 12.3 | 15 |
|                           | T. hemispiralis | 11  | 14  | 12.7| 13 | 0.8 | 6.3  | 15 |
|                           | T. kiaochowensis| 8   | 9   | 8.8 | 9  | 0.5 | 5.2  | 12 |
|                         | T. uruguayensis | T. hemispiralis | T. kiaochowensis |
|-------------------------|-----------------|-----------------|-----------------|
| Shortest kinety in left field, length | 6  8  7.3  7  0.6  8.2  15 | 4  8  5.8  6  1.3  22.8  15 | 3  6  4.6  4  0.9  19.8  15 |
| Shortest kinety in left field, number of kinetids | 3  4  3.6  4  0.5  14.4  12 | 3  4  3.6  4  0.5  14.4  12 | 2  2  2.0  2  0.0  0.0  15 |
| Lateral ciliary field, number of kineties | 9  14  11.9  11  2.3  19.1  15 | 9  17  12.4  12  1.5  12.5  15 | 13  19  15.9  15  1.8  11.5  12 |
| Lateral ciliary field, length of the longest kinety | 9  17  12.4  12  1.5  12.5  15 | 9  17  12.4  12  1.5  12.5  15 | 22  33  28.9  31  4.0  13.7  12 |
| Lateral ciliary field, length of the shortest kinety | 9  14  11.8  12  1.6  13.7  15 | 9  17  12.4  12  1.5  12.5  15 | 22  33  28.9  31  4.0  13.7  12 |
| Kineties in ciliary field, distance to anterior end of cell | T. hemispiralis | T. uruguayensis | T. kiaochowensis |
|                         | 9  14  11.8  12  1.6  13.7  15 | 9  17  12.4  12  1.5  12.5  15 | 13  19  15.9  15  1.8  11.5  12 |
| Adoral zone of membranelles, diameter | T. hemispiralis | T. uruguayensis | T. kiaochowensis |
|                         | 27  57  40.7  40  8.5  20.8  15 | 20  21  17.0  17  1.9  11.1  15 | 16  18  16.3  16  0.5  2.8  12 |
| Collar membranelles, number | T. hemispiralis | T. uruguayensis | T. kiaochowensis |
|                         | 33  52  42.9  42  5.2  12.0  15 | 18  19  18.2  18  0.4  2.3  15 | 16  18  16.3  16  0.5  2.8  12 |
| Buccal membranelle, number | T. hemispiralis | T. uruguayensis | T. kiaochowensis |
|                         | 1  1  1.0  1  0.0  0.0  15 | 1  1  1.0  1  0.0  0.0  15 | 1  1  1.0  1  0.0  0.0  15 |
| Prolonged membranelles, number | T. hemispiralis | T. uruguayensis | T. kiaochowensis |
|                         | 4  5  4.7  5  0.5  10.5  15 | 4  5  4.7  5  0.5  10.5  15 | 3  3  3.0  3  0.0  0.0  12 |
CV, coefficient of variation in %; M, median; Max, maximum; Mean, arithmetic mean; Min, minimum; N, number of specimens examined; SD, standard deviation.

Table 2. DNA sequences obtained in this study.

| Species         | Marker            | Length (bp) | GC content (%) | GenBank accession number |
|-----------------|-------------------|-------------|-----------------|--------------------------|
| *T. hemispiralis* | SSU rDNA         | 1,644       | 47.45           | MT435073                 |
|                 | ITS1-5.8S rDNA-ITS2 | 493         | 46.04           | MT435060                 |
|                 | LSU rDNA          | 1,704       | 51.23           | MT435076                 |
| *T. kiaochowensis* | SSU rDNA        | 1,681       | 47.06           | MT435074                 |
|                 | ITS1-5.8S rDNA-ITS2 | 418         | 45.93           | MT435061                 |
|                 | LSU rDNA          | 1,695       | 50.91           | MT435077                 |
| *T. uruguayensis* | SSU rDNA        | 2,105       | 47.32           | MT435075                 |
|                 | ITS1-5.8S rDNA-ITS2 | 445         | 44.97           | MT435062                 |
|                 | LSU rDNA          | 1,687       | 50.50           | MT435078                 |

Figures
Figure 1

Line drawings of Tintinnopsis hemispiralis in vivo (A, B) and after protargol staining (C–E) (from authors’ own work). A, Lateral view of a representative individual; arrow denotes the ciliary tuft; arrowheads mark the spiral striations on the collar portion of lorica. B, Cell characters; arrow denotes the ciliary tuft; arrowhead shows peduncle. C, D, Ventral (C) and dorsal (D) views of the same specimen, showing ciliary pattern and macronuclear nodules. E, Kinetal map of a morphostatic specimen. BM, buccal membranelle; CM, collar membranelle; DK, dorsal kinety; EM, endoral membrane; LA, lateral ciliary field; LF, left ciliary field; Ma, macronuclear nodule; PCM, prolonged collar membranelle; PK, posterior kinety; RF, right ciliary field; VK, ventral kinety. Scale bars = 75 μm (A, B), 30 μm (C, D).
Figure 2

Photomicrographs of Tintinnopsis hemispiralis in vivo (A–E) and after protargol staining (F–J). A, Lateral view of a representative individual. B, Arrowheads show the spiral striations on lorica. C, Fully extended individual with broken lorica; arrowhead shows the ciliary tuft. D, Lorica of another individual. E, Cell proper that abandoned the lorica; arrowhead shows elongated anterior cilia. F, Ventral kinety and right ciliary fields; arrow shows the ciliary tuft; arrowheads indicate the nodules of the thick argyrophilic fibers. G, Dorsal side, showing the left ciliary field, dorsal kinety, and posterior kinety; arrow marks the ciliary tuft.
Figure 3

Line drawings of Tintinnopsis kiaochowensis in vivo (A, D) and after protargol staining (B, C, E) (from authors' own work). A, Lateral view of a representative individual; arrowheads mark elongated anterior cilia of right and left ciliary field. B, C, Ventral (B) and dorsal (C) views of the same specimen, showing ciliary pattern and macronuclear nodules; arrowhead denotes the left ciliary field. D, Cell features; arrowheads mark elongated anterior cilium of the right and left ciliary fields; arrow shows peduncle. E, Kinetal map of a morphostatic specimen. BM, buccal membranelle; CM, collar membranelle; DK, dorsal kinety; LA, lateral ciliary field; LF, left ciliary field; PCM, prolonged collar membranelle; PK, posterior kinety; RF, right ciliary field; VK, ventral kinety. Scale bars = 50 μm (A), 20 μm (B, C), 40 μm (D).
Figure 4

Photomicrographs of Tintinnopsis kiaochowensis in vivo (A–D) and after protargol staining (E–K). A, Lateral view of a representative individual. B, Different individual showing lorica variation. C, Elongated anterior cilia of the right and left ciliary fields (arrowhead). D, Pressed lorica showing aligned particles. E, F, Arrowheads mark endoral membrane. G, Collar membranelles. H, Ventral side; arrowhead shows the lateral ciliary field. I, Dorsal side of the same specimen as in (H). J, Dorsal (arrowhead) and posterior kinety. K, Subapical view, showing the left ciliary field and dorsal kinety. CM, collar membranelle; DK,
dorsal kinety; LF, left ciliary field; PK, posterior kinety; RF, right ciliary field; VK, ventral kinety. Scale bars = 45 μm (A, B, D), 15 μm (H, I), 20 μm (J).

Figure 5

Line drawings of Tintinnopsis uruguayensis in vivo (A) and after protargol staining (B–D) (from authors’ own work). A, Lateral view of a representative individual. B, C, Ventral (B) and dorsal (C) views of the same specimen, showing ciliary pattern and macronuclear nodules. D, Kinetal map of a morphostatic specimen. BM, buccal membranelle; CM, collar membranelle; DK, dorsal kinety; L, lorica; LA, lateral ciliary field; LF, left ciliary field; PCM, prolonged collar membranelle; PK, posterior kinety; RF, right ciliary field; VK, ventral kinety. Scale bars = 30 μm (A), 15 μm (B, C).
Figure 6

Photomicrographs of Tintinnopsis uruguayensis in vivo (A–D) and after protargol staining (E–J). A, Lateral view of a representative individual. B, Lorica showing a flared collar with jagged rim. C, Lorica with atypical collar rim. D, Cell proper escaped from lorica. E, Ventral side; arrowhead indicates the second short kinety in the right ciliary field. F, Left ciliary field and dorsal kinety. G, Dorsal kinety. H, Prolonged collar membranelles and macronuclear nodules. I, Right ciliary field. J, Lateral view of a middle divider. DK, dorsal kinety; LF, left ciliary field; Ma, macronuclear nodule; OP, oral primordium; PK, posterior kinety; RF, right ciliary field; VK, ventral kinety. Scale bars = 30 μm (A, B), 15 μm (D–K).
Figure 7

Maximum likelihood (ML) tree inferred from SSU rDNA sequences, showing nodal support for ML and Bayesian Inference (BI) analyses. Newly sequenced species are shown in bold. Asterisks (*) reflect disagreements in topology between the BI and ML trees; black circles reflect fully-supported nodes. The scale bar corresponds to 0.05 substitutions per site.
Figure 8

Maximum likelihood (ML) tree inferred from ITS1-5.8S rDNA-ITS2 sequences, showing nodal support for ML and Bayesian Inference (BI) analyses. Newly sequenced species are shown in bold. Asterisks (*) reflect disagreements in topology between the BI and ML trees; black circles reflect fully-supported nodes. The scale bar corresponds to 0.05 substitutions per site.
Figure 9

Maximum likelihood (ML) tree inferred from LSU rDNA sequences, showing nodal support for ML and Bayesian Inference (BI) analyses. Newly sequenced species are shown in bold. Asterisks (*) reflect disagreements in topology between the BI and ML trees; black circles reflect fully-supported nodes. The scale bar corresponds to 0.1 substitutions per site.
Figure 10

Sampling sites. A, Map of China with sample sites (yellow circles), downloaded from the open-access website: www.osgeo.cn. B, C, Photographs of Meng Bay and coast of Qingdao, respectively. Note: The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.

Supplementary Files

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- Supplementarymaterials.pdf