

Observations of Apparent Lorica Variability in *Salpingacantha* (Ciliophora: Tintinnida) in the Northern Pacific and Arctic Oceans

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Abstract. Species identifications of tintinnid ciliates are based on characteristics of the lorica housing the ciliate cell. Molecular characterization of tintinnid ciliates has revealed the occurrence of cryptic species, genetically distinct forms with similar loricas, as well as polymorphic species in which genetically identical forms have distinct loricas. Among this latter category may be forms currently recognized as species of the genus *Salpingacantha*, erected by Kofoid and Campbell for forms originally described as varieties of species of *Salpingella* with ‘teeth’ on the margin of the lorica oral opening. Some workers have expressed the opinion that the genus is artificial and notably a recent molecular study suggested that *Salpingella* and *Salpingacantha* are likely synonyms. In this short communication we report on finding morphologically distinct loricas (containing ciliate cells) of *Salpingacantha* in single samples from stations in the North Pacific and Arctic Oceans in 2010, 2015 and 2016. We found up to 5 distinct *Salpingacantha* forms co-occurring, often with abundant *Salpingella acuminata* populations suggesting that some *Salpingacantha* species may be varieties of *Salpingella acuminata*. While we lack genetic data needed for definitive proof of polymorphism, here we document remarkable gradual gradients in morphology suggestive of polymorphism.

Keywords: microzooplankton, plankton, polymorphism, tintinnida.

INTRODUCTION

The genus *Salpingacantha* was created by Kofoid and Campbell (1929) separating forms previously considered as varieties of *Salpingella* which had ‘teeth’ on the oral margin (*S. undata*, *S. simplex*, *S. unguiculata*) as well 4 ‘new’ forms, with teeth more or less evident

on the oral margins, that they described from the Agassiz Expedition (*S. ampla*, *S. crenulata*, *S. exilis*, *S. perca*). In their revision of 1939, Kofoid and Campbell state “Closely resembles *Salpingella* but differs from it in that the oral margin is toothed or has at least the first phases in the formation of teeth, as in *perca*, in which they are mere minute marginal extensions of longitudinal folds”. With regard to *Salpingacantha simplex*, their description 1929 states “... collar scarcely flaring, without definite teeth, its margin undulating...”. Thus although ‘teeth’ on the oral margin supposedly distinguish the genus *Salpingacantha*, exactly what constitutes ‘teeth’ is unclear. Furthermore, the appearance (or

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not) of ‘teeth’ could depend on small changes in the orientation of a specimen or small deformations in the oral margin.

A recent revision of tintinnid taxonomy based on molecular data suggests that *Salpingacantha* is likely a synonym of *Salpingella* (Santoferrara *et al.* 2017). The observation was based on findings that the SSU rDNA sequences of two forms corresponding with the morphologies described for *Salpingacantha undata* and *S. unguiculata* were intermingled among sequences attributed to *Salpingella acuminata*. Their opinion of synonymy is not new. Over the years various authors have opined that given the overall similarities in lorica morphology, *Salpingacantha* is probably a synonym of *Salpingella* (e.g., Travers and Travers 1973; Cordeiro and Sassi 1997; Alder 1999). In fact, Cordeiro and Sassi (1997) provided a series of micrographs illustrating a remarkable gradient of morphologies ranging from *Salpingella acuminata* to *Salpingacantha undata*. However, most of the specimens were empty lorica and different lorica were from different samples. It is important to distinguish empty lorica from those containing ciliate cells. Empty lorica are known to persist in the water column, can become degraded or distorted, and have wider geographic distributions than the living cells (Kato and Taniguchi 1993). Empty lorica are then unreliable indicators of living ciliates. Evidence of the co-occurrence of live forms of distinct *Salpingacantha* appears to be lacking and provided the impetus for this short report. Here we report on the co-occurrence of apparent morphological variants in *Salpingacantha* in samples gathered in the summer from 2010 to 2016 in the North Pacific and Arctic Oceans.

METHODS AND MATERIALS

Samples were collected from onboard the Korean Research Icebreaker *Araon* in August/early September from 2010 to 2016. Sampling dates and station locations of samples considered in this report are given in the supplementary data file. Plankton net tows were performed to assess microplankton community composition. Net tows were made using a 20- μ m plankton net of 0.45 m diameter usually towed from 100 m depth to the surface (water column depth permitting). Net tow material was fixed by standard methods (6% Bouin’s or 2% Lugol’s, final concentration). Aliquots were examined in settling chambers. Specimens correctly oriented for diagnosis were enumerated and photographed using an Olympus IX71 microscope equipped with DIC optics and a DP71 camera interfaced with a PC running Olympus BioCell image analysis software with scale calibration (Olympus France SAS). In the text the sample origins are indicated in parentheses in bold indicating year and station.

The location coordinates and sampling dates appear in the supplementary data file. Tintinnid “species” identifications based on lorica morphology refer to the designations found in the monographs of Kofoid and Campbell (1929, 1939).

RESULTS AND DISCUSSION

On several occasions we found a variety of different *Salpingacantha* forms within single samples. *Salpingacantha* were generally, but not always, found in presence of *Salpingella acuminata*. In one sample (**2015 St 5**) we found 5 morpho-types representing a clear gradient from *Salpingella acuminata* to *Salpingacantha undata* (Fig. 1). As the length of the outward sloping teeth alters greatly lorica opening diameter (LOD), the forms differed considerably in LOD, ranging from 18 μ m for the crenulated oral margin form (Fig. 1c) to the ‘4-toothed’ with an LOD of 27 μ m (Fig. 1f). Notably in two samples *Salpingacantha* were found in the absence of *Salpingella*. Figure 2 shows 5 different morpho-types found in a sample (**2016 St 29**) in which no typical *Salpingella acuminata* were found.

Different morpho-types of *Salpingacantha* dominated different samples with no apparent pattern. The numbers and dimensions of the different morpho-types in individual samples is given in the supplementary data file. Among specimens of *Salpingacantha* there was relatively little variability in overall length but distinct differences in the diameter and shape of the lorica oral opening. We suggest that the co-occurrence with *Salpingella acuminata* and the large gradient in architecture of the lorica oral opening of *Salpingacantha* raises the possibility that “species” of *Salpingacantha* may be varieties of *Salpingella* species.

Salpingacantha and *Salpingella* are “cosmopolitan” genera (Pierce and Turner 1993), both widely (and co-incidentally) distributed (Dolan and Pierce 2013, Fig. 10.3). However, *Salpingella* species are found much more commonly than *Salpingacantha*. A quick search using *Google Scholar* (<https://scholar.google.com>) for publications containing the term ‘*Salpingella*’ yields about 500 publications whereas the term ‘*Salpingacantha*’ returns about 50. Most (but not all) reports of the occurrence of *Salpingacantha* species also report the occurrence of *Salpingella*.

There has never been a consensus with regard to whether or not *Salpingacantha* are morpho-types of *Salpingella* or distinct species. Early on Jørgensen, in creating the genus *Salpingella* (Jørgensen 1924), con-

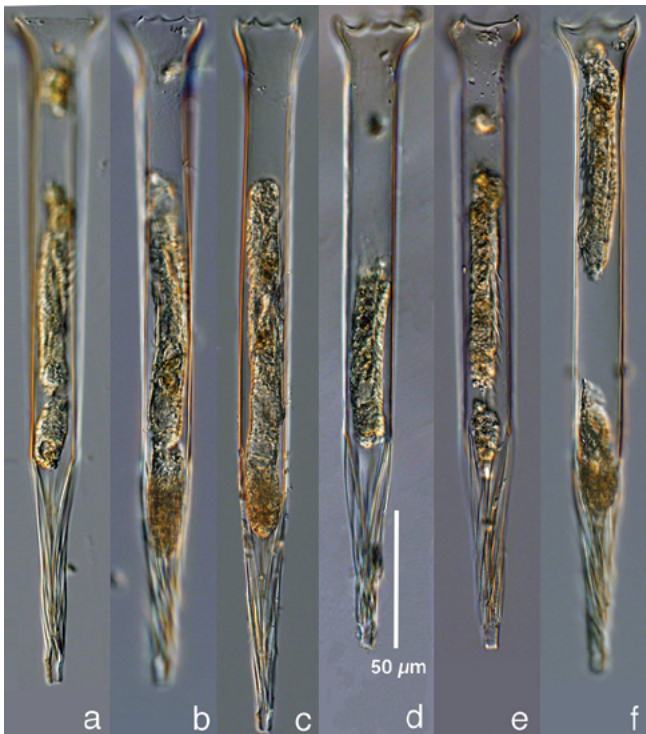


Fig. 1. Varieties of *Salpingacantha* from a 2015 sample in the Chukchi Sea (2015 St 5). In the sample, typical *Salpingella acuminata* (“a”) were dominant with 70 cells found for the 15 *Salpingacantha* cells encountered. The “b” most closely corresponds with *S. crenulata*, “c” and “d” with *S. perca*, “e” with *S. unguiculata* and “f” with *S. ampla*. Note that form “e”, if rotated 45°, could easily be mistaken for *Salpingella acuminata*.

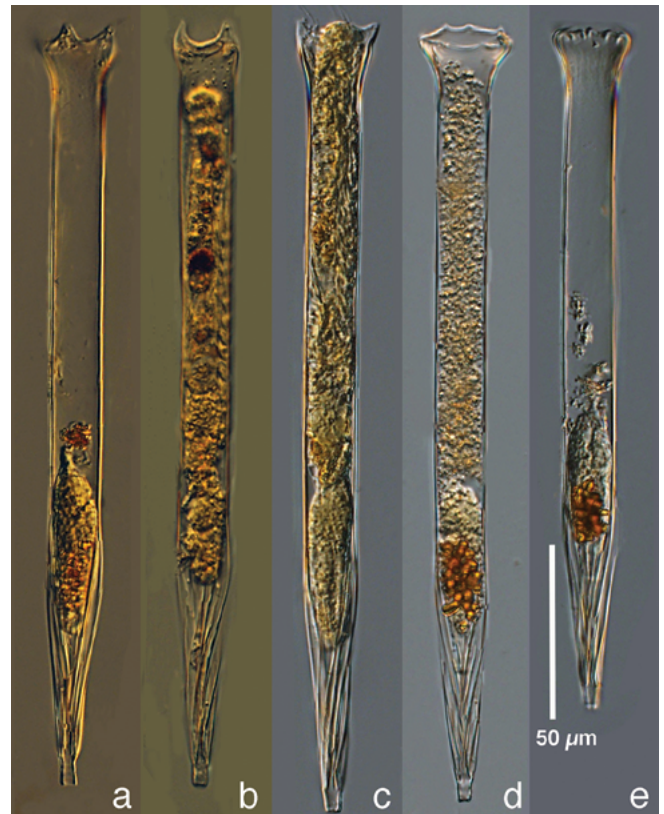


Fig. 2. Varieties of *Salpingacantha* from a 2016 sample in the Chukchi Sea (2016 St 29). In the sample no typical *Salpingella acuminata* were found. In the conspectus of Kofoid and Campbell (1929) the form “a” most closely resembles *S. ampla*, “b” *S. unguiculata*, “c” *S. simplex*, “d” *S. perca*, and “e” *S. crenulata*. Note that form “c” if rotated 45° could easily be mistaken for *Salpingella acuminata*.

tended that *Salpingella acuminata* with “teeth” were an artifact, perhaps mounting media distorts a relatively flexible oral margin; he disputed Brandt’s designation of toothed forms as separate species (Brandt 1906). Kofoid and Campbell (1939: 394) discounted Jörgensen’s view that the ‘teeth’ were an aberration or an artifact. However, they admitted that in their samples *Salpingella* usually occurred with *Salpingacantha* and furthermore that *Salpingacantha* usually occurred as a small number of individuals.

It is perhaps significant that for Kofoid and Campbell, the forms they distinguished as species of *Salpingacantha* represented an evolutionary series of toothed oral margin beginning with *S. perca* and culminating in *S. undata* as shown in Fig. 3 (Kofoid and Campbell 1939: 394). Kofoid was an avowed proponent of orthogenesis (see Kofoid 1930) and it is plausible to

assume he was strongly pre-disposed to seeing order and directionality in diverse morphologies. Admittedly, we can be assumed to be pre-disposed to assuming that tintinnid species can exhibit more or less distinct morphologies. We have previously found polymorphism in *Cymatocylis* (Kim *et al.* 2012) with different morpho-types not clearly associated with environmental conditions but rather multiple morpho-types found when population density is high (Dolan *et al.* 2013). Obviously, a definitive answer with regard to *Salpingacantha* awaits single-cell sequencing to diagnose the genetic identity of all the diverse morpho-types we and others (Cordeiro and Sassi 1997) have encountered.

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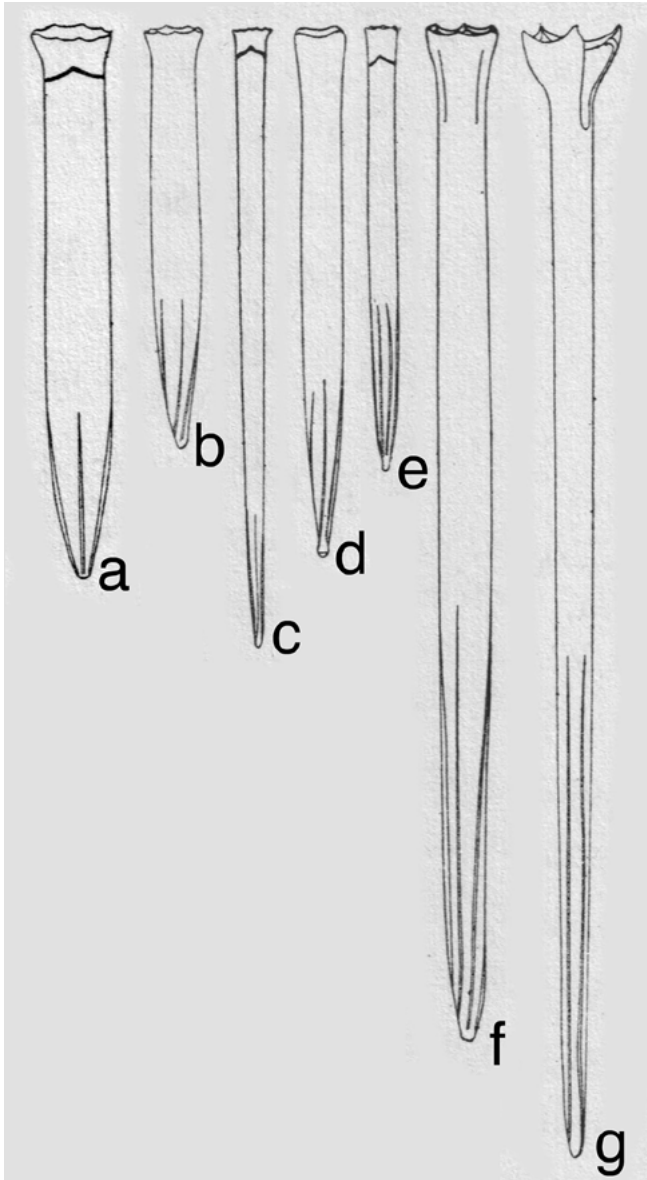


Fig. 3. The illustrations of *Salpingacantha* species from Kofoid and Campbell 1929 arranged as the evolutionary series given in Kofoid & Campbell 1939 in which *S. perca* “presents the earliest phase in evolution of toothed oral margin”: a. *S. perca*, b. *S. crenulata*, c. *S. exilis*, d. *S. simplex*, e. *S. unguiculata*, f. *S. ampla*, and g. *S. undata*. Note that *S. perca*, *S. exilis*, and *S. unguiculata* are depicted with a diaphragm apparatus similar to those of *Salpingella* (Agatha 2010).

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