

# First records of *Amphidoma languida* and *Azadinium dexteroporum* (Amphidomataceae, Dinophyceae) from the Irminger Sea off Iceland

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*Species of dinophycean Amphidomataceae are producers of phycotoxins classified as azaspiracids. We provide the first records of two of their constituent species, Amphidoma languida and Azadinium dexteroporum, for the Irminger Sea off Iceland. Morphological examination and molecular characterization, including uncorrected mean pairwise distances between sequences of the Internal Transcribed Spacer (ITS), doubtlessly assigned the sub-Arctic strain 2A11 to the reference of Amphidoma languida. Strain 2A11 produced AZA-38 and AZA-39, corresponding to the toxin profile described for the type material. The sub-Arctic isolate 1D12 differed significantly in terms of ITS genetic distance ( $p = 0.04$ ) from a Mediterranean Azadinium dexteroporum strain, but our morphological analysis did not reveal any major or stable diagnostic traits between the reference strain of Azadinium dexteroporum and the new strain described here. In contrast to the Mediterranean strain of Azadinium dexteroporum, the sub-Arctic strain 1D12 did not produce any known azaspiracids. The new records of Amphidoma languida and Azadinium dexteroporum from the Irminger Sea imply an important range extension of the species, formerly known from the northern and eastern Atlantic (Amphidoma languida) and from the Mediterranean area (Azadinium dexteroporum) only. Together with three new species of Azadinium recently described from the same expedition, the results clearly show that the biodiversity of the Amphidomataceae in the sub-Arctic is remarkably large.*

**Keywords:** *Azadinium*, *Amphidoma*, azaspiracids, Irminger Sea

Submitted 15 January 2015; accepted 17 August 2015

## INTRODUCTION

It is predicted that climate change will have dramatic consequences for the (sub-)Arctic region, including increased mean temperature and rapid decline of ice cover (Comiso *et al.*, 2008; Screen & Simmonds, 2012). These environmental alterations have the potential to cause shifts and changes in plankton communities (Wassmann *et al.*, 2011) and, in particular, may expand the distribution of harmful algae bloom (HAB) species into or within the Arctic Sea (Hallegraeff, 2010). However, to fully evaluate any potential future changes in species distribution and community composition, basic information about the occurrence and distribution of species (past and present) is needed.

Starting in the first half of the 19th Century (e.g. Ehrenberg, 1843), there is a long history on plankton surveys in the Arctic seas. Expeditions during the late 19th and early 20th Century (e.g. Cleve, 1873; Cleve & Grunow, 1880; Brandt & Apstein, 1908; Lebour, 1925; Grontved & Seidenfaden, 1938) resulted

in extensive lists of plankton species, which remain the basis for contemporary taxonomic research in the Arctic. However, most of these studies focused on diatoms and larger dinophytes because of their using plankton nets for collection and light microscopy for observation. With respect to planktonic protists, the recent assessment of the pan-Arctic biodiversity (Poulin *et al.*, 2011) consequently reveals a striking paucity of information about smaller plankton species (<20  $\mu\text{m}$ ), accounting for less than 20% of the currently known species from the region.

A bias in the composition of species lists is also obvious for planktonic dinophytes: among about 200 species present in the Arctic Ocean compiled by Okolodkov & Dodge (1996), more than 40% belong to three generic lineages (*Dinophysis*, *Protoperidinium*, *Tripes*) of large and conspicuous thecate species. Thus, the biodiversity and distribution of smaller and less conspicuous dinophytes is not fully explored at this moment in time, as it is illustrated by, for example, *Azadinium* (Amphidomataceae). Driven by the specific search for the planktonic source of azaspiracids (AZAs) (Krock *et al.*, 2009), the taxon was discovered in 2009 with the description of *Azadinium spinosum* Elbrächter & Tillmann. This small and inconspicuous species is the nano-planktonic source of AZA-1, -2, and -33 (Tillmann *et al.*,

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2009, 2012a). Since then, knowledge about the diversity of *Azadinium* has increased rapidly and today ten species are described (Tillmann *et al.*, 2014). At least two other species of *Azadinium* also produce AZAs (i.e. *Azadinium poporum* Tillmann & Elbrächter; Krock *et al.*, 2012; *Azadinium dexteroporum* Percopo et Zingone; Percopo *et al.*, 2013). With the description of *Amphidoma languida* Tillmann, Salas & Elbrächter, another nanoplanktonic AZA-producing organism of a divergent (but closely related) lineage of Amphidomataceae was recently identified (Tillmann *et al.*, 2012b).

Most species of *Azadinium* have been initially described from the North Sea (Tillmann *et al.*, 2009, 2010, 2011) but today, multiple records from various parts of the world are available (Akselman & Negri, 2012; Hernández-Becerril *et al.*, 2012; Potvin *et al.*, 2012; Gu *et al.*, 2013; Luo *et al.*, 2013; Percopo *et al.*, 2013). However, knowledge about Amphidomataceae, and the presence of AZAs, in the (sub-)Arctic is still scarce (Poulin *et al.*, 2011). *Azadinium caudatum* (Halldal) Nézan & Chomérat, the largest and most conspicuous species of *Azadinium*, has initially been described (as *Amphidoma caudata*) from the northern Norwegian coast in winter (Halldal, 1953). In addition, a species described as *Gonyaulax parva* Ramsfjell from the central Norwegian Sea towards Iceland (Ramsfjell, 1959) almost certainly is also a species of *Azadinium* (see discussion in Tillmann *et al.*, 2014), while another species from the Canadian Arctic (Holmes, 1956; Bérard-Therriault *et al.*, 1999) described as *Gonyaulax gracilis* J. Schiller probably also refers to a species of *Azadinium* (see discussion in Tillmann *et al.*, 2014).

In 2012, a research cruise to the North Atlantic Ocean near Greenland and Iceland was carried out, and a number of strains of small Amphidomataceae were isolated. In total, cultures of seven different strains were established. Five of these strains representing three new species of *Azadinium* (*Azadinium trinitatum* Tillmann & Nézan, *Azadinium cuneatum* Tillmann & Nézan, *Azadinium concinnum* Tillmann & Nézan) have been described elsewhere (Tillmann *et al.*, 2014). Here, we present the detailed identification and the AZA toxin profile of the two other strains, which represent

the first sub-Arctic records of *Amphidoma languida* (initially described from the Irish coast: Tillmann *et al.*, 2012b) and *Azadinium dexteroporum* (previously known from the Mediterranean only: Percopo *et al.*, 2013).

## MATERIAL AND METHODS

### Isolation and cultivation

Two strains designated as 2A11 and 1D12 were established from water samples collected between Greenland and Iceland during a cruise aboard the research vessel 'Maria S. Merian' in August 2012 (Figure 1). Strain 1D12 was isolated from the Irminger Sea (Station 526: 64°45.71'N 29°56.74'W), whereas strain 2A11 was isolated off the north-west coast of Iceland (Station 532: 65°27.00'N 24°39.00'W). Isolation and establishment of clonal strains was performed as described by Tillmann *et al.* (2014). Established strains were routinely grown at both 10°C and 15°C in a natural seawater medium prepared with sterile-filtered (0.2 µm VacuCap filters, Pall Life Sciences; Dreieich, Germany) Antarctic seawater (salinity: 34 psu, pH adjusted to 8.0) and enriched with 1/10 strength K-medium (Keller *et al.*, 1987), slightly modified by omitting additive NH<sub>4</sub><sup>+</sup> ions. Strains were grown in 250 ml plastic culture flasks at 15°C under a photon flux density of 50 µmol m<sup>-2</sup> s<sup>-1</sup> on a 16:8 h light:dark photocycle.

### Microscopy

Light microscopy (LM), including epifluorescence microscopy of 4'-6-diamidino-2-phenylindole (DAPI) stained nuclei, and scanning electron microscopy (SEM) were performed as described by Tillmann *et al.* (2014). Cell length and width were measured at 1000x microscopic magnification using Zeiss Axiovision software (Zeiss, Germany) and freshly fixed cells (formaldehyde, final concentration 1%) of cultures growing at 15°C.

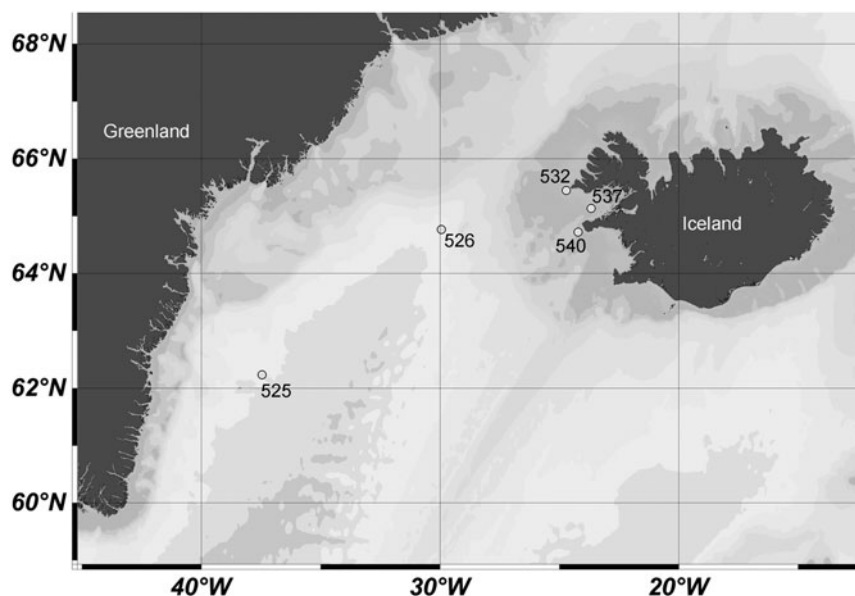


Fig. 1. Selected sampling localities of the 'Maria S. Merian' expedition 2012.

## Toxin analyses

For each harvest, cell density was determined by settling lugol-fixed samples and counting >800 cells under an inverted microscope (Axiovert 200M, Zeiss; Göttingen, Germany). Dense cultures (densities ranging from ca.  $20\text{--}80 \times 10^3$  cells ml<sup>-1</sup>) were harvested in  $4 \times 50$  ml centrifuge tubes by centrifugation (Eppendorf 5810R; Hamburg, Germany) at 3220 g for 10 min. Each four pellets from one harvest were combined in an Eppendorf microtube and again centrifuged (Eppendorf 5415R; Hamburg, Germany) at 16,000 g for 5 min and stored frozen ( $-20^\circ\text{C}$ ) until use. The growth and harvest procedure was repeated several times to yield a total number of at least  $1.4 \times 10^8$  cells. The total volume and number of cells harvested was 4.4 l,  $1.48 \times 10^8$  cells, and 11.0 l,  $1.45 \times 10^8$  cells, for *Amphidoma languida* and *Azadinium dexteroporum*, respectively.

All harvests of each strain were combined in 2 ml methanol and homogenized with a sonotrode (70 s, 70 cycles, 100% power; Sonoplus HD 2070, Bandelin, Berlin, Germany). Homogenates were centrifuged ( $15^\circ\text{C}$ , 3220 g, 15 min; Eppendorf 5810 R, Hamburg, Germany). Supernatants were collected and pellets re-extracted twice with 1 ml methanol, each. Combined extracts were reduced in a rotary evaporator (Büchi, Konstanz, Germany) at reduced pressure and  $40^\circ\text{C}$  water bath temperature to a volume <0.5 ml and then taken up in acetone to a final volume of 1 ml. The extracts were transferred to a  $0.45\text{-}\mu\text{m}$  pore-size spin-filter (Millipore Ultrafree, Eschborn, Germany) and centrifuged at 800 g for 30 s (Eppendorf 5415R), with the resulting filtrate being transferred into a liquid chromatography (LC) autosampler vial for LC coupled to tandem mass spectrometry (LC-MS/MS) analysis.

To analyse AZAs, samples were analysed by LC-MS/MS according to the methods described in detail by Tillmann *et al.* (2009). Selected reaction monitoring experiments were carried out in positive ion mode by selecting the transitions given in Table 1. In addition, precursor ion experiments were performed as described in detail by Tillmann *et al.* (2014).

## Molecular phylogenetic analysis

DNA isolation, rRNA fragment amplification and sequencing followed standard methods described in Tillmann *et al.* (2014). Sequences of the new strains were included in an alignment covering the known diversity of the Amphidomataceae. Phylogenetic methods were the same as specified in Tillmann *et al.* (2014). Uncorrected mean pairwise distances (proportion: p-distances) between sequences of the Internal Transcribed Spacer (ITS) were calculated with PAUP\* version 4.0b10 (Swofford, 2002). The sequences were also inspected for possible compensatory base changes that occur when substitutions take place in pairing regions of the molecule folded into a secondary structure (Gottschling & Plötner, 2004; Thornhill & Lord, 2010; Kremp *et al.*, 2014).

## RESULTS

### Morphological characterization

Motile cells determined as *Azadinium* sp. or *Amphidoma* sp. were observed in concentrated whole water samples at a

**Table 1.** Mass transitions m/z ( $Q_1 > Q_3$  mass) and their respective AZAs.

Mass transition	AZA	Collision energy (V)
716 > 698	AZA-33	40
816 > 798	AZA-34, AZA-39	40
816 > 348	AZA-39	70
828 > 810	AZA-3	40
828 > 658	AZA-3	70
830 > 812	AZA-35, AZA-38	40
830 > 348	AZA-38	70
842 > 824	AZA-1, AZA-6, AZA-40	40
842 > 672	AZA-1	70
842 > 348	AZA-40	70
844 > 826	AZA-4, AZA-5	40
846 > 828	AZA-37	40
846 > 348	AZA-37	70
854 > 846	AZA-41	40
854 > 670	AZA-41	70
856 > 838	AZA-2	40
856 > 672	AZA-2	70
858 > 840	AZA-7, AZA-8, AZA-9, AZA-10, AZA-36	40
858 > 348	AZA-36	70
860 > 842	Undescribed	40
872 > 854	AZA-11, AZA-12	40

number of stations in the Irminger Sea between Greenland and Iceland, and around the north-west coast of Iceland (Figure 1). From this cruise, a total of seven clonal strains were established, two of which were determined as *Amphidoma languida* (2A11) and *Azadinium dexteroporum* (1D12), respectively, based on morphology and rRNA sequence comparison.

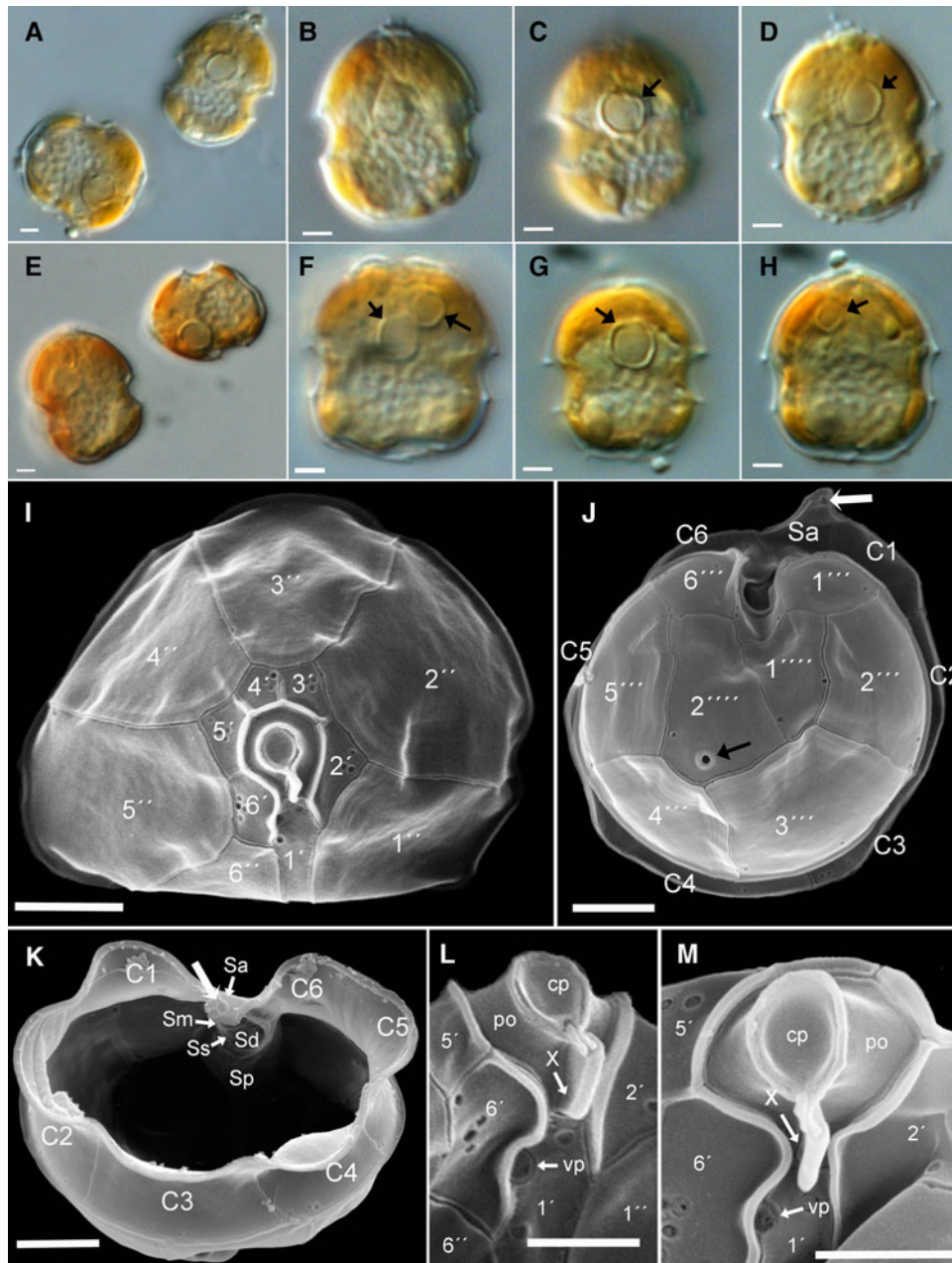
### AMPHIDOMA LANGUIDA 2A11

Cells of the strain 2A11 were easy to identify at low magnification LM because of their characteristic swimming pattern. Cells regularly concentrated at the bottom of the cultivation vessel and exhibited a conspicuous behaviour, in that they normally moved very slowly, and suddenly and irregularly performed jumps. These jumps occurred when cells approached the bottom of the cultivation vessel, or were stimulated by short vibrations of the cultivation vessel.

Cells of *Amphidoma languida* (strain 2A11) were ovoid to slightly elliptical (Figure 2A–D), with a conical episome and a distinctly pointed apex. Mean size of recently formaldehyde-fixed cells was  $13.8 \pm 0.9 \mu\text{m}$  (min 11.8–max 16.3,  $N = 112$ ) in length and  $11.8 \pm 1.0 \mu\text{m}$  (min 9.6–max 15.1,  $N = 112$ ) in width, resulting in a mean length/width ratio of 1.2. The episome was slightly larger than the hemispherical hyposome, which ended in a pointed antapex (Figure 2D). A large-lobed chloroplast engrossed the entire cell (Figure 2E), usually with a single large pyrenoid centrally located in the episome (Figure 2A, C–E). Rarely, cells with two pyrenoids were observed (Figure 2F). As quantified once, 35 of 600 cells (5.8%) had two pyrenoids. If two pyrenoids were present, then mostly they were both located in the episome (rarely one of the two pyrenoids was located in the hyposome) and could be of different sizes (Figure 2G, H). The large and spherical to ovoid nucleus was located in the posterior part of the cell (Figure 2A–H).

The plate formula was determined by SEM as Po, cp, X, 6', oa, 6'', 6C, 5(?)S, 6''', 2''''.





**Fig. 2.** *Amphidoma languida* (strain 2A11). (A–H) LM of formalin fixed cells. (A–E) General size and shape. Note the presence of a large pyrenoid in the episome (arrows in C, D). (F–H) Variations in pyrenoid. (F) A broad cell with two pyrenoids (arrows) in the episome. (G, H) Two focal planes of one cell indicating the presence of a large and a small pyrenoid (arrows) in the episome. (I–M) SEM of different cells showing plate pattern of the epitheca (I), hypotheca (J), and of the cingular and sulcal plates in interior view (Sa, anterior sulcal plate; Sm, median sulcal plate; Sd, right sulcal plate; Ss, left sulcal plate; Sp, posterior sulcal plate) (K). (L, M) Cell apex (po, pore plate; cp, cover plate; X, X-plate; vp, ventral pore). Black arrow in J indicates the large antapical pore, white arrows in J and K the roundish depression on plate Sa. Scale bars: (A–K) = 2  $\mu\text{m}$ ; (L, M) = 1  $\mu\text{m}$ .

specific thecal plates, the location of a ventral pore on plate 1' and the location of a large antapical pore on plate 2''', as well as the arrangement of small pores on the plates are shown in Figure 2I–M. On the anterior sulcal plate, a roundish depression was detected at the anterior tip (white arrows in Figure 2J, K). Unlike a pore, the depression did not seem to pass through the thecal plate.

#### AZADINIUM DEXTEROPORUM 1D12

The second strain 1D12 was clearly different from *Amphidoma*, even at low magnification LM, because of its

conspicuously smaller size and a different swimming pattern. Cells of strain 1D12 performed sudden jumps as well interrupting a generally calm swimming style, but occasionally they covered longer distances in a faster and jerky mode.

The mean cell size of strain 1D12 was  $10.5 \pm 0.6 \mu\text{m}$  (min 9.1–max 12.3, N = 172) in length and  $7.3 \pm 0.6 \mu\text{m}$  (min 6.0–max 8.9, N = 172) in width, with a mean length/width ratio of 1.4. The cingulum was very broad and deeply excavated (Figure 3A–H). A distinct antapical spine was visible in LM (Figure 3D, G–H). A presumably single chloroplast

was parietally arranged and lobed (Figure 3E, F). In cells of a dense culture, the chloroplast was often restricted to the episome, although fluorescence microscopy clearly indicated autofluorescence in both epi- and hyposome (Figure 3I–K). A relatively small pyrenoid visible by its starch cup was present (Figure 3C, E, G–H). In addition to the pyrenoid, cells may have a number of large granular structures mainly located in the hyposome, which differed from the pyrenoids in the absence of a clear starch shield covering them (Figure 3H). Among 600 cells examined in LM at 1000x magnification, two cells were found with two small pyrenoids in the episome, whereas all other cells had a single pyrenoid (sometimes difficult to be identified) consistently located in the episome. A spherical or slightly elongated nucleus was located in the posterior part of the cell. Occasionally (presumably at first stage of cell division), the nucleus distinctly elongated and almost completely filled the cell (Figure 3I–K).

The Kofoidian plate pattern of Po, cp, X, 4', 3a, 6'', 6C, 5(?)S, 6''', 2'''' (Figure 4) conformed with the most common plate pattern of the whole *Azadinium* genus. The three lateral and dorsal apical plates were small. The sutures of the dorsally located plate 3' to its neighbouring apical plates were very short so that the epithelial intercalary plates were almost adjacent to the pore plate (Figure 4D, I). Three symmetrically arranged intercalary plates were located dorsally. The outer intercalary plates each bore a group of small pores, whereas plate 2a invariably was free of pores (Figure 4B–E). As the most abundant arrangement, the distinctly smaller central intercalary plate 2a was quadrangular and almost symmetrically located above plate 3'' (Figure 4B, D). A penta-configuration (i.e. plate 2a was pentagonal) was at times present with plate 2a in contact with 3'' and 4'' (Figure 4C, E).

An oval- to teardrop-shaped apical pore was located in the centre of the pore plate (Po) and was connected through a finger-like protrusion with the small X-plate (Figure 4I–M). When seen from inside the cell, the X-plate was small and ovoid (Figure 4N) and was posteriorly tangent but not invading the first apical plate.

As the most characteristic feature, the ventral pore (vp) of *Azadinium dexteroporum* was located at the distal end of the

more-or-less elongated right side of the asymmetric pore plate. In the strain 1D12, a large variability in shape of the pore plate and, thus, in the position of the ventral pore (Figure 4I–M) was found. The vp was displaced posteriorly in varying degrees at the tip of the asymmetric right side of the pore plate but rarely, it was located almost next to the X-plate on the slightly elongated right part of the pore plate (Figure 4I). The pore plate was surrounded by a raised proximal rim of the four apical plates all along its posterior margin. This rim continued ventrally along the sutures of 1' and 4' bordering the vp (Figure 4I–M), although the rim around the vp varied in thickness and was partially missing occasionally (Figure 4J). The vp itself had a thick outer rim and ranged in outer diameter from 0.26 to 0.46  $\mu\text{m}$  (mean:  $0.36 \pm 0.04 \mu\text{m}$ ,  $N = 21$ ).

The wide cingulum was descending and displaced by half of its width. Each of the cingular plates exhibited a few scattered pores. On plate C<sub>1</sub>, and rarely also on plate C<sub>5</sub>, pores occasionally were enriched in conspicuously arranged fields comprising up to 20 pores (Figure 4F, G). The sulcus comprised five main plates (Figure 4H) in the arrangement typical for *Azadinium*, with the left sulcal plate (Ss) running along the line from plate C<sub>1</sub> to C<sub>6</sub>. Two small plates (the median sulcal plate (Sm) and the right sulcal plate (Sd)) formed a concave central pocket. An additional structure was occasionally visible above Sm and Sd (Figure 4H). However, it could not be verified whether this represented an additional sulcal platelet, or an internal outgrowth of plate C<sub>6</sub> extending to both central sulcal plates. On the hypotheca, the larger of the two antapical plates (i.e. plate 2''') exhibited a small spine typically accompanied by a few small pores (Figure 4F).

Desmoschisis, as a type of cell division, is described here for the first time for *A. dexteroporum* (Figure 5). The parental theca was shared between the two sister cells, and dividing cells retained their motility throughout the complete mitotic and cytokinetic process. Prior to division, cells increased mainly in width (Figure 5A) and later, plates were separated along a well-defined oblique fission line dividing an anterior-sinistral part from a posterior-dextral part (Figure 5B–F, schematized in Figure 5G–H). Along the fission line, the anterior-

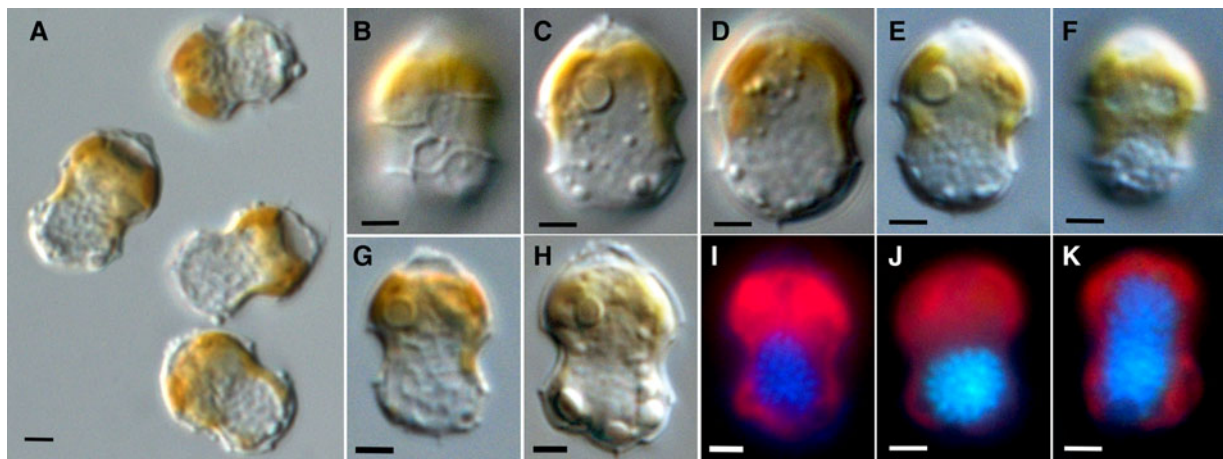
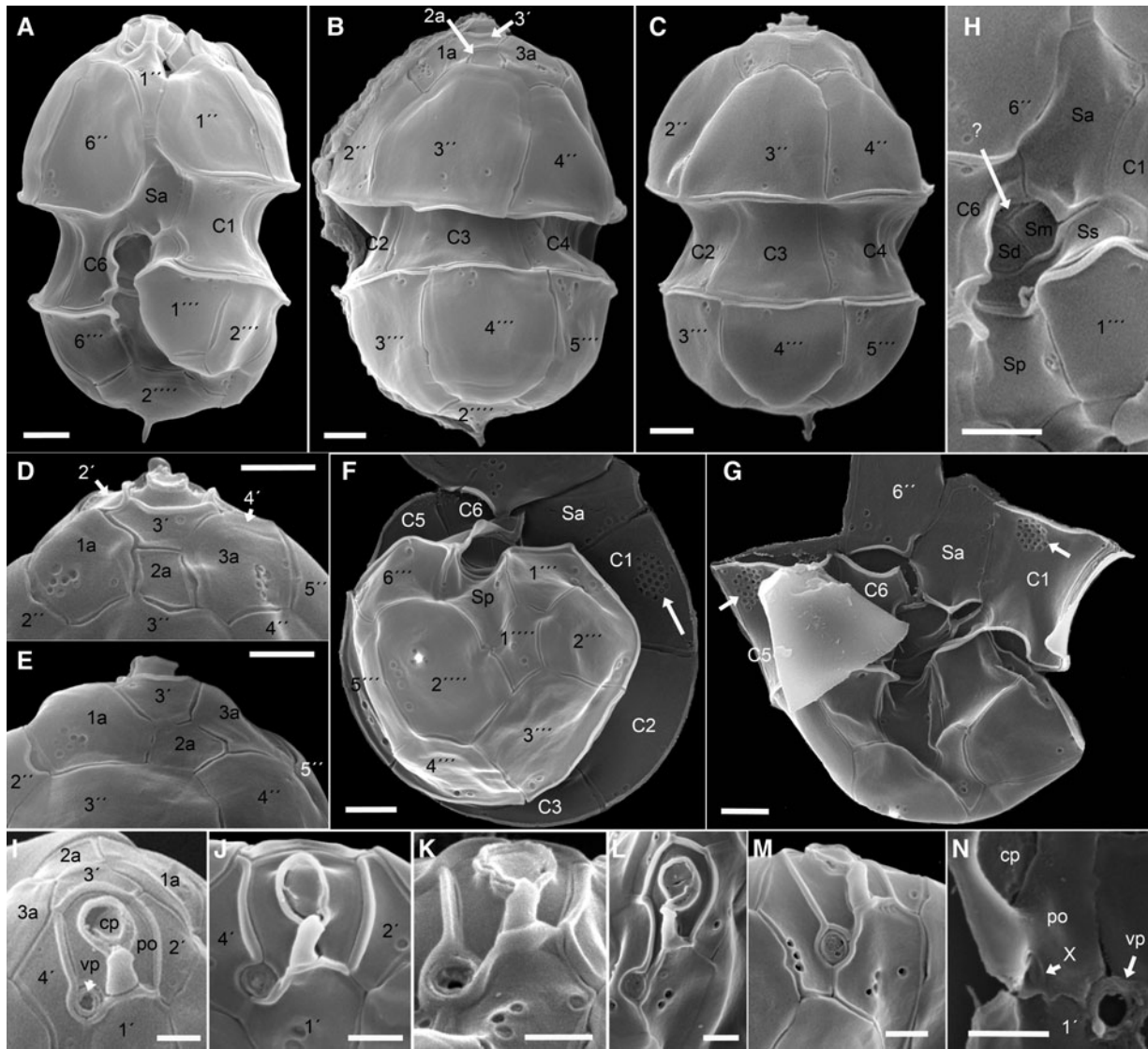


Fig. 3. *Azadinium dexteroporum* (strain 1D12). LM of formaldehyde-fixed cells. (A–H) General size and shape. Note the presence of a large pyrenoid in the episome (C, E, G, H) and the presence of an antapical spine (D, G, H). (E, F) Two focal planes of the same cell to illustrate the lobed chloroplast. (H) Cell with a pyrenoid and additional large grains of presumably storage material in the episome. (I–K) Formalin-fixed cells stained with DAPI under UV excitation showing nucleus and chloroplast shape and position. Scale bars = 2  $\mu\text{m}$ .



**Fig. 4.** *Azadinium dexteroporum* (strain 1D12). SEM of different cells. (A) Ventral view. (B, C) Dorsal view. (D, E) Detailed dorsal view showing plate 2a in quadra- (D) or penta-configuration (E). (F) Antapical view showing hypothecal and cingular plates. Note a large field of scattered pores on the first cingular plate (arrow). (G) Field of pores on C1 and C5 (arrows). (H) Detailed view of the sulcal plates. Note the additional structure (?) visible above Sm and Sd (Sa, anterior sulcal plate; Sm, median sulcal plate; Sd, right sulcal plate; Ss, left sulcal plate; Sp, posterior sulcal plate). (I–N) Detailed views of the apical area (po, pore plate; cp, cover plate; vp, ventral pore). (N) Internal view of the apical area (X, X-plate). Scale bars: (A–G) = 1  $\mu$ m; (I–N) = 0.5  $\mu$ m.

sinistral daughter cell (light shaded) inherited all apical plates (including the apical pore complex), the three anterior intercalary plates, the first two precingular plates, the anterior sulcal plate (Sa) and the first four postcingular plates. The postero-dextral daughter cell (darker shading) received the remaining precingular plates (i.e. 3''–6''), the posterior sulcal plate (Sp), the fifth and sixth postcingular plates, and the two antapical plates.

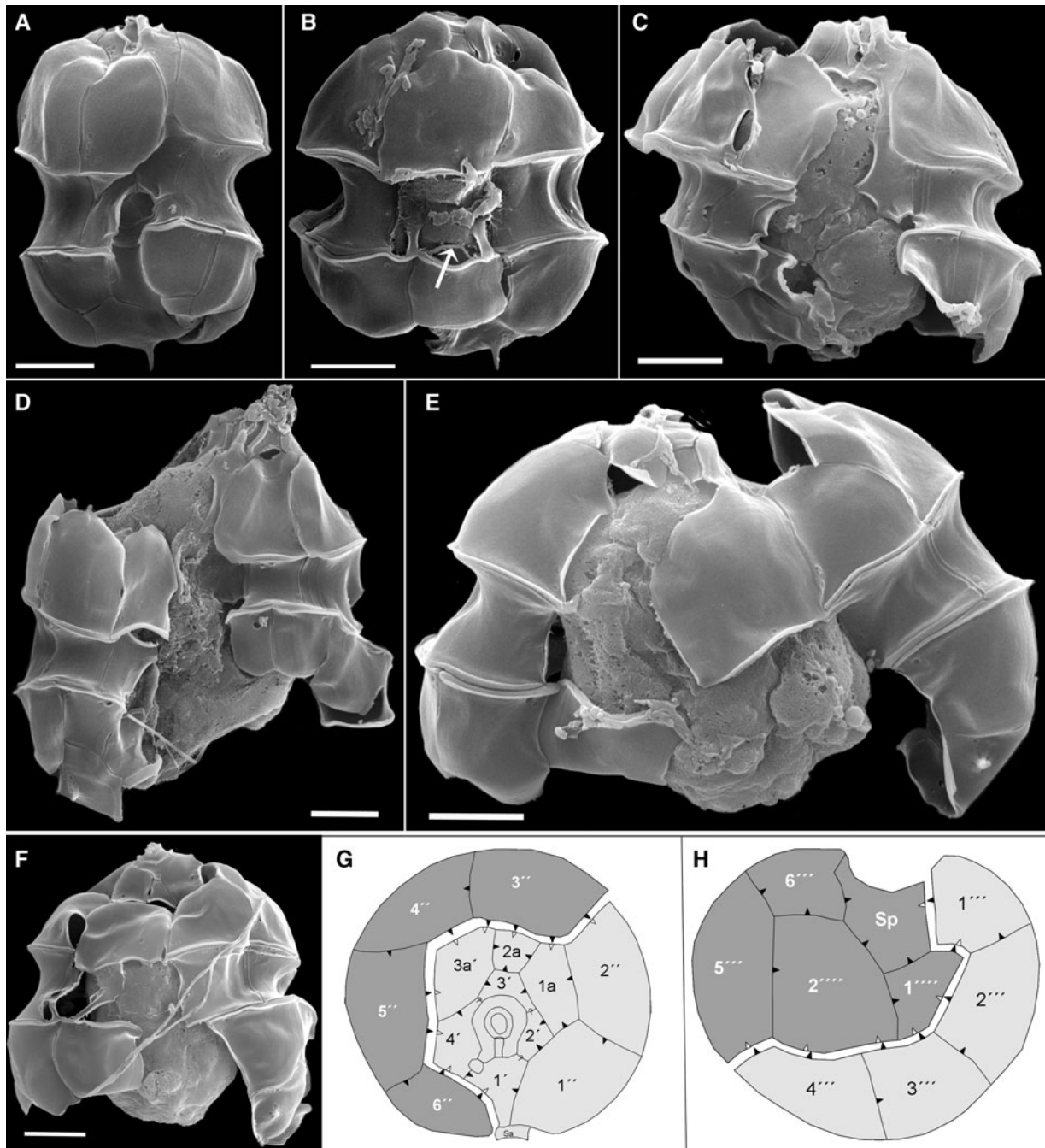
Interior views of thecate cells with slightly disarranged plates allowed an identification of the overlap patterns for most plates (Figure 5G–H). Plate 2a was overlapped by all adjacent plates, and plate C6 was overlapped by the central sulcal plate Sa. As keystone plates (i.e. plates that overlap all adjacent plates) of *A. dexteroporum*, plates 3'', C3 and 4''' for the precingular, the cingular and the postcingular series were identified, respectively. The course of the fission line followed this plate overlap pattern. In the epitheca, all plates of the postero-dextral daughter cell overlapped plates of the

anterio-sinistral daughter cell (Figure 5G). In the hypotheca, plates of the anterio-sinistral half overlapped plates of the postero-dextral half with one exception: the first antapical plate overlapped plate 1''' of the anterio-sinistral daughter cell (Figure 5H).

## Phylogenetics

New rRNA sequences were submitted to GenBank and are available as entries KR362880–KR362890. The SSU + ITS + LSU alignment was 4609 bp long and comprised 1813 parsimony informative sites (39%, mean of 11.62 per operational taxonomic unit). The Amphidomataceae, including species of *Amphidoma* and *Azadinium*, were monophyletic (99LBS, 1.00BPP; Figure 6). The new strains were reliably retrieved (100LBS, 1.00BPP) with *Amphidoma languida* (strain 2A11) and *Azadinium dexteroporum* (strain 1D12), respectively.



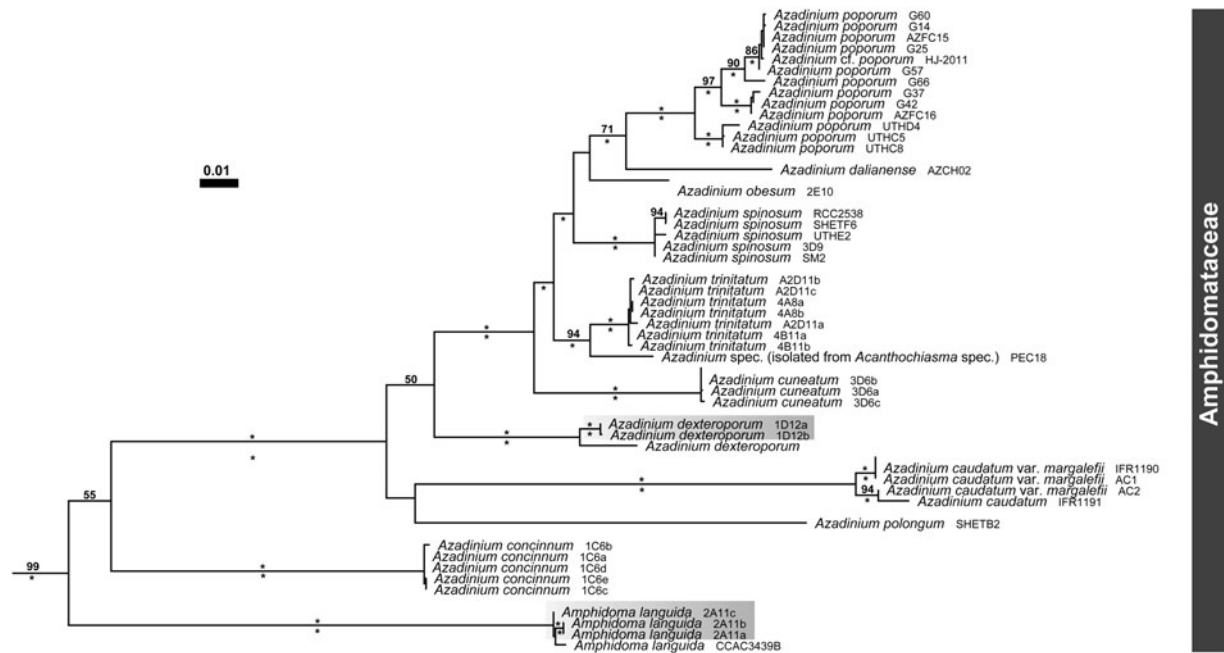


**Fig. 5.** *Azadinium dexteroporum* (strain 1D12). SEM of different cells. (A) Pre-division cell with a broadened cell width. (B–F) Cells in different stages of cell division in ventral (C, D) or dorsal (B, E, F) view (arrow in (B): indication of dissolution of cingular plate C<sub>3</sub>). (G, H) Schematic view of fission line of epitheca (G) and hypotheca (H) separating the anterio-sinistral daughter cell (light shading) and the postero-dextral daughter cell (darker shading); arrowheads indicate direction of plate overlap, with black symbols at the fission line indicating plate margins overlapping the plate with the corresponding white symbols. ? = direction of plate overlap unclear. Scale bars = 2  $\mu\text{m}$ .

Within strains and species, ITS pairwise distances were low (Table 2), specifically  $\sim 0.01$  between the Irish strain SM1 of *Amphidoma languida* and strain 2A11, and  $\sim 0.04$  between the Mediterranean strain SZNB-848 of *Azadinium dexteroporum* and strain 1D12, respectively. All LSU sequences of *Amphidoma languida* obtained are identical, whereas the LSU sequence of SZNB-848 exhibited seven different positions in comparison to those (identical) of 1D12. Compensatory base changes could not be stated, also not for the more divergent strains of *Azadinium dexteroporum*.

### Azaspiracids

Strain 2A11 (*Amphidoma languida*) showed AZA cell quotas of 0.8 and 1.5  $\text{fg cell}^{-1}$  for AZA-38 and AZA-39, respectively. In the cell pellet of strain 1D12 (*Azadinium dexteroporum*), known AZAs were not detected. The limit of detection of AZAs was calculated as 0.007  $\text{fg cell}^{-1}$  for AZA-1. For both strains, precursor scans of the characteristic AZA fragments  $m/z$  348 and  $m/z$  362 did not reveal any indication of the presence of yet undescribed variants.



**Fig. 6.** Maximum likelihood tree ( $-\ln = 72,424.15$ ) of 48 operational taxonomic units assigned to the Amphidomataceae (cut-off from Tillmann *et al.*, 2014, with the outgroup truncated), as inferred from a MAFFT-generated rRNA nucleotide alignment spanning the SSU, ITS and LSU (1813 parsimony informative positions). Branch lengths are drawn to scale, with the scale bar indicating the number of nucleotide substitutions per site. Numbers on branches are statistical support values for the clusters to the right of them (above: ML bootstrap support values, values under 50 are not shown; below: Bayesian posterior probabilities, values under 0.90 are not shown), and asterisks indicate maximal support values. The new sub-Arctic strains are highlighted.

## DISCUSSION

The data now available leave no doubt that strain 2A11 represents the first record of *Amphidoma languida* from the sub-Arctic. The conspicuous swimming behaviour has been previously described and documented for *Amphidoma languida* (Tillmann *et al.*, 2012b) and thus, this strain has been expected to represent this particular species since first observation. The specific designation is confirmed by detailed LM and SEM studies: plate formula, arrangement, size and shape of specific thecal plates, the location of a vp on plate 1' and the location of a large antapical pore on plate 2''', as well as the arrangement of small pores scattered over the plates, correspond to the original description of *Amphidoma languida* (Tillmann *et al.*, 2012b). The roundish depression on the anterior tip of the anterior sulcal plate has not been explicitly noted for the type material, but it is visible in the original material of strain SM1 (Tillmann *et al.*, 2012a). However, such a depression is not always obvious (or visible) and thus cannot be regarded as a consistent morphological trait.

The sub-Arctic records of *Amphidoma languida* represent a remarkable range extension of the species. Formerly, it has

been described from the Irish coast (Salas *et al.*, 2011) and most probably has been depicted as *Amphidoma* sp. by Lewis & Dodge (1990) from material collected in the eastern North Atlantic. In contrast to the shallow coasts of Ireland and the sub-Arctic near Iceland, cells most likely determinable as *Amphidoma languida* have been observed in SEM from a sample collected at the open West-Indian Ocean as well ( $5^{\circ}02.4'S$   $45^{\circ}16.8'E$ ; Consuelo Carbonell-Moore, personal communication), indicating a global distribution and broad ecological tolerance of this species.

Molecular phylogenetic data are in agreement with the morphological conclusion with respect to *Amphidoma languida*. Sequences of the strain 2A11 are almost identical to the reference of *Amphidoma languida*, as inferred from the low ITS pairwise distances, which are in the usual range for the categories 'intra-genomic' and 'individuals of same species' (Litaker *et al.*, 2007). On the other hand, ITS pairwise distances are larger between the reference sequence of *Azadinium dexteroporum* and the sequences obtained from the new strain described here. They are even larger than between geographic variants of *Azadinium poporum* ( $p < 0.02$ : Gu *et al.*, 2013) or between the two morphotypes of

**Table 2.** Genetic pairwise p-distances between ITS regions from *Amphidoma languida* (first three columns) and *Azadinium dexteroporum*.

	<i>Amphidoma languida</i> strain SM1	2A11 sample 13-368 ITS clone 2, 9	<i>Azadinium dexteroporum</i> SZN-B848	1D12 Sample 12-316
2A11 Sample 13-368 ITS clone 2	0.01353	0.00000	—	—
2A11 Sample 13-368 ITS clone 9	0.01353	—	—	—
2A11 Sample 13-368 ITS clone 10	0.00846	0.00839	—	—
1D12 Sample 12-316	—	—	0.03994	—
1D12 Sample 13-133	—	—	0.04111	0.00000



*Azadinium caudatum* var. *caudatum* and *Azadinium caudatum* var. *margalefii* (Rampi) Nézan & Chomérat ( $p = 0.023$ ; Nézan *et al.*, 2012). A  $p$ -distance of 0.04 is set as the threshold for distinguishing species (Litaker *et al.*, 2007), although fixed values can always be not more than approximation for biological phenomena.

Basic differences in the autecology of the Mediterranean and the sub-Arctic strains of *Azadinium dexteroporum* seem to be present. In the Mediterranean, the species has been identified at water temperatures between 17.8°C and 23.8°C, and the strain in cultivation has not survived at temperatures below 18°C (Percopo *et al.*, 2013). In contrast, our strain 1D12 has been isolated from water with a temperature of 8°C and is now routinely grown in the lab at 10°C and 15°C, respectively. However, our morphological analysis has not revealed diagnostic traits between the reference of *Azadinium dexteroporum* and the new strains described here, with thecal plate details corresponding to the type material (Percopo *et al.*, 2013).

However, some minor morphological differences between the two strains of *Azadinium dexteroporum* described in detail so far are worthy of discussion. In sub-Arctic strain 1D12, the position of the vp is slightly variable but generally closer to the apical pore plate than it has been reported for the Mediterranean (type) material. Furthermore, we have never observed a pronounced concavity of the median intercalary plate 2a, which has been highlighted as a peculiar feature of the Mediterranean (type) strain (Percopo *et al.*, 2013). In strain 1D12, the median intercalary plate is either tetragonal (i.e. in contact with four other epithecal plates) or pentagonal (and in contact with five epithecal plates), but the latter shape has not been reported for Mediterranean *Azadinium dexteroporum*. Division mode, fission line and plate overlap pattern reported here for the first time for *Azadinium dexteroporum* exactly resemble the state for the type *Azadinium spinosum* as described by Tillmann & Elbrächter (2010, 2013).

Species delimitation in unicellular protists such as the dinophytes is challenging for multiple reasons, such as complex determination procedures (mostly under the microscope only) and uncertainty of morphological diagnostic traits (frequently at high intraspecific variability). It also remains an open question at this moment in time whether the minute differences between the Mediterranean and sub-Arctic strains of *Azadinium dexteroporum* in terms of morphology, molecular phylogenetics and ecology are consistent and are an expression of biological speciation, or are to be interpreted as intraspecific variability. More comparative studies (both in the field and using cultivated material), and particularly breeding experiments between different strains, are necessary to resolve this problem. Cryptic speciation is not restricted to the Amphidomataceae, but is a great taxonomic challenge also in other dinophyte lineages such as *Alexandrium* (Kremp *et al.*, 2014) and *Scrippsiella* (Zinßmeister *et al.*, 2011; Kretschmann *et al.*, 2014, 2015).

The azaspiracid profile of strain 2A11 determined as *Amphidoma languida* with the presence of AZA-38 and AZA-39 is identical to the (type) strain SM1 (Krock *et al.*, 2012), and toxin cell quotas (fg per cell range) are similar for both strains. The absence of AZAs in strain 1D12 determined as *Azadinium dexteroporum* is in contrast to the results of Percopo *et al.* (2013) reporting from the presence of AZAs in the Mediterranean strain. However, it is known

that toxin production can be variable among strains of a single species. *Azadinium poporum* has been first described as a non-toxicogenic species (Tillmann *et al.*, 2011), but produces several different, novel AZAs (Krock *et al.*, 2012) with a high variability within strains. Among a total of 22 strains of *Azadinium poporum* analysed so far, four strains without any detectable AZAs were found (Gu *et al.*, 2013; Krock *et al.*, 2014). Thus, toxin analyses of more strains are needed to evaluate if the presence/absence of AZAs is a consistent phenotypic trait with the potential to differentiate between Mediterranean and sub-Arctic strains of *Azadinium dexteroporum*.

The new records of *Amphidoma languida* and *Azadinium dexteroporum*, together with the description of three new species of *Azadinium* (Tillmann *et al.*, 2014) from the same expedition to the Irminger Sea, clearly show that the diversity of the Amphidomataceae in the sub-Arctic is large, especially since our presented findings are based on a single cruise and a limited number of stations. The results of studies like the present are needed to complement and broaden a pan-Arctic biodiversity inventory (Poulin *et al.*, 2011).

## ACKNOWLEDGEMENTS

Thanks to Captain Bergmann and the FS Maria S. Merian crew for their assistance and support for the collection of field material. We greatly acknowledge the help of Wolfgang Drebing (AWI) for analyses of azaspiracids. Financial support was provided by the PACES research program of the Alfred Wegener Institute as part of the Helmholtz Foundation initiative in Earth and Environment.

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