**Taxonomic re-investigation and geochemical characterization of Reid’s (1974) species of *Spiniferites* from holotype and topotype material**

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**Abstract**

The genus *Spiniferites* currently encompasses 142 dinoflagellate cyst species. Some *Spiniferites* species are difficult to identify because of an incomplete or doubtful description, and/or substandard iconography. This study re-describes and re-illustrates the *Spiniferites* holotypes first described by Reid in 1974. It also discusses topotype material from surface sediments recovered from British estuaries, and attempts to provide further constraints on the classification of species in this genus using the geochemical characterization of their cyst walls. Reid described four new *Spiniferites* species: *Spiniferites belerius*, *Spiniferites delicatus*, *Spiniferites elongatus* and *Spiniferites lazus*. New photomicrographs are presented here for the holotypes of *S. delicatus* and *S. elongatus*, and additional morphological observations based on newly processed topotype material are given. The geochemical characterization of the *Spiniferites* cyst walls showed overall consistency with a carbohydrate-based dinosporin. However, variability in the dinosporins suggests that, in this genus, the cyst wall composition may be species-specific. Analysis of the characteristic spectral regions for unclassified *Spiniferites* species showed that, in some cases, it may be possible to constrain the likely species affinity using the cyst wall chemistry. However, in most cases, the morphologically unspeciated cysts did not show sufficient similarity to an identified species’ cyst wall chemistry to be more conclusive. This could either reflect an intermediate species that cannot be clearly characterized using morphology or dinosporin composition, or it represents a completely different species. In either case, both the morphological and geochemical evaluations highlight the difficulties in classifying species of this genus unequivocally.

**Keywords:** dinoflagellate cyst, redescription, taxonomy, geochemistry, cyst wall composition, FTIR, dinosporin.

**1. Introduction**

The genus *Spiniferites* Mantell 1850, currently encompassing 142 species (Fensome et al., 2008), is renowned for the difficulty in distinguishing different species, often with species morphology intergrading between each other (Mertens et al. this volume). Furthermore, the cyst-theca relationship of many species remains unclear (e.g. Rochon et al. 2009).

Reid (1972, 1974, 1977) studied dinoflagellate cyst assemblages from estuaries around the British Isles (Figure 1). The assemblages were recovered from surface sediments and included 11 species belonging to the genus *Spiniferites* (Mantell 1850). Four species were considered new to science and formally described in Reid (1974): *Spiniferites belerius*, *Spiniferites delicatus*, *Spiniferites elongatus* and *Spiniferites lazus*. The other *Spiniferites* species recovered and documented by Reid (1974) included *Spiniferites bentorii* (Rossignol 1964) Wall & Dale 1970, *Spiniferites bulloideus* (Deflandre & Cookson 1955) Sarjeant 1970, *Spiniferites hyperacanthus* (Deflandre & Cookson 1955) Cookson & Eisenack 1974, *Spiniferites membranaceus* (Rossignol 1964) Sarjeant 1970, *Spiniferites mirabilis* (Rossignol 1964) Sarjeant 1970, *Spiniferites pachydermus* (Rossignol 1964) Reid 1974 and *Spiniferites ramuliferus* (Deflandre 1937) Reid 1974. Reid (1974) transferred the latter species from *Achomosphaera*, and it is now considered synonymous with *Achomosphaera ramosasimilis* (see Mertens et al. this volume). In this study, the holotypes of two of the four new *Spiniferites* species defined by Reid (1974), *Spiniferites delicatus* and *Spiniferites elongatus*,were re-examined and re-photographed. Newly processed sediment samples from topotype localities allowed further observations to elucidate the morphology of the species and document them with new photomicrographs.

 In addition to the taxonomic re-evaluation, the composition of several *Spiniferites* species’ cyst walls, including three of the species described by Reid (1974), *Spiniferites belerius*, *Spiniferites delicatus*, and *Spiniferites elongatus*, were measured and analyzed using micro-Fourier transform infrared (FTIR) spectroscopy. Micro-FTIR is a widely used technique for the evaluation of biopolymers (Olcott-Marshall & Marshall 2015, and references therein) and has been successfully used in the geochemical analysis of both fossil (e.g., Versteegh et al. 2007, Bogus et al. 2012) and recent (e.g., Kokinos et al. 1998; Versteegh et al. 2012; Bogus et al. 2014; Mertens et al. 2015a, 2015b, this volume) dinoflagellate cysts.

Dinoflagellate cyst walls are composed of dinosporin, suggested to be a complex carbohydrate-based biomacromolecule. Dinosporin was initially described as such based on FTIR and pyrolysis gas chromatography-mass spectrometry of *Lingulodinium machaerophorum* from culture and surface sediments (Versteegh et al. 2012); additional analyses suggested that considerable differences in composition between species of the same genus are possible (Bogus et al. 2012). Thus, it appears phylogeny could be a factor contributing to dinosporin compositional differences. However, this suggestion is limited by the fact that species from different genera, environmental conditions (surface waters and post-depositional) and regions have primarily been compared (e.g., Bogus et al. 2014). These other factors seemingly influence the dinosporin composition (e.g., Bogus et al. 2014; Mertens et al. 2015a, 2015b, this volume), a finding consistent with the fact that dinoflagellate cyst morphology as well can vary with differing environmental conditions (e.g., Ellegaard 2000; Lewis et al. 2001; Ellegaard et al. 2002; Zonneveld & Susek 2007; Mertens et al. 2009).

By comparing the cyst wall chemistry of *Spiniferites* species from the same location (Dee Estuary), likely deposited at about the same time, we can constrain most of these uncertainties. This allowed us to investigate the extent to which a particular dinosporin composition may be species specific. Our results were then used in an attempt to constrain the speciation of specimens that could not be visually assigned because of unclear morphology.

**2. Material and methods**

***2.1. Observation of holotypes and topotypes***

The holotypes of *Spiniferites belerius*, *Spiniferites delicatus*, *Spiniferites elongatus* and *Spiniferites lazus* are stored in the repository at the Micropalaeontology Laboratory, Geology Department of the University of Sheffield (UK) under catalogue numbers 69K1 1331.568(4), 12 ABSL1 1290.192(5), 140 K3 1210.352(12) and 73K1 1365.256(2), respectively. These specimens were re-examined by P.R.G. during summer 2014.

Topotype material was sampled in the Dee, Caernarvon and Conwy Marina estuaries (Figure 1). The saltmarsh surface sediment was carefully hand-collected in a 50 ml plastic tube that was covered with black tape. All samples were then kept in the dark, at 4°C. Additional samples were investigated from previously studied surface samples from the Celtic Sea (location details in Marret et al. 2004) and surface samples from Britanny (Vilaine Estuary, location details in Mertens et al. 2009). The palynological processing of the topotype and additional samples followed the procedure described by e.g. Pospelova et al. (2010). Samples were treated with 10% hydrochloric acid (HCl) at room temperature to remove carbonates. After rinsing with distilled water, samples were treated with hydrofluoric acid (HF, 48-50%, 2 days) to remove silicates, followed by a second 10% HCl treatment to remove precipitated fluorosilicates. The samples were rinsed again and sonicated up to 30 seconds and the residue was sieved and retained on a nylon mesh screen (15 µm). One or two drops of the residue were mounted on a slide with glycerine jelly and covered with a cover slip.

The holotypes were photographed at the University of Sheffield with an Infinity 1 camera mounted on a Meiji MT 5300H microscope, while the specimens from the topotype material were photographed at a 100× magnification with an MRc5 camera mounted on a Zeiss Axio Imager A1 at Ghent University and with an Olympus DP72 camera mounted on a BX41 microscope at the Station de Biologie Marine (Ifremer, Concarneau). For scanning electron microscope (SEM) observations, single cysts were picked from the residue and mounted on a glass slide or were filtered using polycarbonate membrane filters (Millipore, Billerica, MA, USA, GTTP Isopore, 0.22 μm pore size), sputter coated with gold, and examined using a JEOL6400 SEM at Ghent University and a Zeiss SIGMA300 Gemini field emission SEM at the Station de Biologie Marine (Ifremer, Concarneau).

The holotypes of *Spiniferites delicatus* and *Spiniferites elongatus* were successfully identified in the palynological collections at the University of Sheffield. The holotypes of *Spiniferites belerius* and *Spiniferites lazus* were not found due to the poor condition of the slides. All of the *Spiniferites* species identified by Reid (1974), except *Spiniferites bentorii*, *Spiniferites hyperacanthus* and *Spiniferites pachydermus*, were found in the newly processed samples from topotype localities.

Additional observations, for illustrative purposes, of '*Spiniferites ramuliferus'* were made on residues from the type stratum and locality of *Spiniferites coniconcavus* in the Pliocene Verrebroek Dock section in Belgium (processing details in De Schepper et al. 2004).

***2.2. Geochemical analysis of cyst wall chemistry***

The palynological residues from topotype material (Dee Estuary) were then treated for geochemical analysis. They were sonicated and rinsed three times with organic solvents (methanol and dichloromethane) and water to remove polar and apolar compounds that might have adhered to the outside of the cyst walls. Visually clean individual cysts were manually isolated, placed on an Au-coated mirror and dried. Specimens were analyzed with a Bruker Hyperion 2000 microscope coupled to a Bruker Vertex 80v FTIR spectrometer at the Department of Solid State Sciences (Ghent University). Magnification of the microscope was set at 15x and the aperture adapted to the size of the measured specimen. The combination of detector (liquid N2 cooled MCT detector), source (Globar) and beamsplitter (KBr) settings restrict the infrared spectral range examined to ~7000–600 cm-1, from these spectra the 4000–650 cm-1 range was selected for analysis in this study. All presented spectra were collected in reflection mode, at a resolution of 2 cm-1 and averaged over 100 scans.

Data were analyzed with the OPUS software. The presented absorbance spectra were obtained after background (direct reflection on the Au mirror) subtraction and baseline correction (rubberband correction method using polynomes and 5 iterations). Residual CO2 absorption traces at ~2350 cm-1, a result of incomplete N2 gas purging of the microscope, have also been removed. Functional groups were identified based on comparison with published literature. Relative strengths of the main IR band regions within a specimen’s spectrum were calculated by integrating the area under the peaks between defined band limits to allow comparisons between spectra. The ranges of these IR bands are: (I) 3010–2775 cm-1, (II) 1850–1500 cm-1, (III) 1500–1185 cm-1, (IV) 1185–860 cm-1.

**3. Systematic paleontology**

Division DINOFLAGELLATA (Bütschli 1885) Fensome et al.1993, emend. Adl et al. 2005

Class DINOPHYCEAE Pascher 1914

Subclass PERIDINIPHYCIDAE Fensome et al.1993

Order GONYAULACALES Taylor 1980

Suborder Gonyaulacineae autonym

Family Gonyaulacaceae Lindemann 1928

Subfamily Gonyaulacoideae autonym

Genus *Spiniferites* Mantell 1850, emend. Sarjeant 1970

*Spiniferites belerius* Reid 1974

Plate 3, Figures 1–12.

**Synonymy.** non *Spiniferites belerius* in Harland 1977, p. 98–99, Plate 1, Figures 7–10, Plate 2, figures 7–10, 16–21, 25–57 [these images correspond to *Spiniferites membranaceus*; Harland (1979) also reports *Spiniferites belerius* from the Pliocene and Pleistocene of the Bay of Biscay, but does not provide photographs; the confirmation of a correct identification in that paper can thus not be given].

**Dimensions.** Reid (1974) – holotype: central body: 35–29 µm; length processes: 10 µm. Range: length central body: 35–42 µm; width central body: 28–37 µm; depth 33–37 µm. Maximum process length: 7–10 µm; maximum posterior process length: 10–15 µm. Number of specimens measured: 15.

Our measurements on topotype material from the Dee, Conwy Marina and Caernarvon estuaries – central body length 32 (34.8) 38 µm; width 27.6 (30.2) 32 µm; process length 6.4 (8.3) 11 µm. Number of specimens measured: 7. These new measurements are comparable with those described by Reid (1974) although somewhat in the lower part of the range. This is similar to the measurements reported by Harland (1977).

**Original description (Reid 1974, p. 597).** “Test oval, always widest at the girdle; at times constricted towards the poles to give a rounded diamond shape with a clear apical node. The thin smooth surfaced wall is ornamented by gonal trifurcate processes formed from sutural septae. Septae form the high antapical ‘trumpet’ shaped process which characterizes the species and may also form high membranous flanges. Process tips are trifurcate with recurved bifurcate tips. A relatively narrow 6–8 µm girdle is displaced by one to three times its own width.”

**Additional observations.** Typically, these cysts are the smallest *Spiniferites* species in the assemblages. The cysts are ovoid with a very finely granular wall (both endophragm and periphragm) (Plate 3, Figure 12). A small apical boss is present (Plate 3, Figure 10). Sutures reveal a tabulation typical for the genus of 4´, 6´´, ?c, ?s, 6´´´, 1p, 1´´´´ (not reported by Reid 1974). The suture between 1´ and 4´ is faint but visible, while 6´´ is triangular. The sulcal plates are faintly discernible, but we could not observe all of them. The archeopyle corresponds to plate 3´´. The processes are rigid and hollow, and exclusively gonal. The distal ends of the processes are typically trifurcate, and each furcation terminates in bifurcate tips, which can be elongated or truncated (Plate 3, Figure 6). The bases of the processes are wide (7.1 µm on average). Reid (1974) defined *Spiniferites belerius* as characterized by an antapical trumpet-shaped process located at the junction of the 1´´ and 2´´´ plates (i.e. an antapical position). We assume Reid meant the large box-like process we observed at the junction of plates 1´´´´ and 1p (thus left of the sulcus) (Plate 3, Figure 3). We suggest it is formed by the fusion of two adjacent processes.

**Comparison.** *Spiniferites belerius* resembles *Spiniferites coniconcavus* De Schepper et al. (2004) from the Pliocene of Belgium, but the latter lacks an apical boss (Pl. 10, Figures 1–3). *Spiniferites belerius* differs from ‘‘*Spiniferites bulloideus* sensu Wall 1965’’ in that *S. belerius* has an apical boss. Its differences with *Spiniferites* *bentorii* are the larger size, the pear-shaped body, the pronounced apical boss and the occurrence of fenestrate bases in *Spiniferites bentorii*.At this time the presence/absence of an apical boss is considered to vary interspecifically and intraspecifically and there is no conclusive evidence for its taxonomic importance (Mertens et al., this volume).

**Remarks.** The holotype was not located in the original slide, possibly because of the poor condition of the slide. The observed specimens were recorded from the Dee and Caernarvon estuaries and the Conwy Marina.

**Occurrence.** From the base of the Pliocene to Recent. Specimens recorded in older Neogene sediments are mentioned as *Spiniferites* cf. *belerius* (Londeix et al. this volume).

*Spiniferites delicatus* Reid 1974

Plate 1, Figures 1–16 (holotype), Plate 4, Figures 1–9.

**Synonymy.** None.

**Dimensions.** Reid (1974) – holotype: central body: 47 x 49 µm; length processes: 21 µm, width girdle: 7 µm. Range: central body: 40 × 35 to 60 × 54 µm. Maximum height processes: 29 µm; width girdle: 6–9 µm. Number of specimens measured: 18. Our measurements on topotype material from the Irish Sea – length 36.8 (41.5) 50.8 µm; width 40 (42) 46 µm. Number of specimens measured: 5.

**Original description (Reid 1974, p.601)**. “The cyst wall is thick, with radial ‘columellae’ or fibres in the endophragm and with a much thinner granular periphragm. The surface is microgranular to microreticulate. Processes are supported by thin skeletal rods which trifurcate and then bifurcate in the process tips. These processes are usually connected by high granular membranous flanges which vary greatly in their development in individual specimens. An apical node or low boss may be found at the head of 1´ and 4´. In most specimens the sutural flanges are of equal height over the whole test, but a few may have high antapical flanges surrounding 1´´´´. The girdle is sinistral and is displaced by three times its width. The sulcus which is orientated posterior/anteriorly [*sic*]. It has the following sulcal plates: PS, IS, LS, RA and AS. The tabulation is typical for the genus with an apical series of four plates and a precingular archeopyle 3´´ which is reduced.”

**Additional observations.** The central body of the cyst is spherical to ovoidal. Our observations and measurements indicate that the central body of the specimens from the topotype material are mostly spherical, and somewhat smaller compared to the measurements of Reid (1974). The inner and outer walls are granular, closely appressed except at the base of the processes. The sutural crests are high, particularly on the antapical and cingular sides. The crests are granular and connect the exclusively gonal processes, which are also granular. The distal ends of the processes are consistently petaloid in plan view. Both processes and crests have faintly granular surfaces. The girdle is sinistral and is displaced by three times its width.

**Comparison.** *Spiniferites ristingensis* Head 2007 (Plate 9, Figures 1–8) has similar processes, but the wall appears to be thinner with a tegillum that forms blisters and lower sutural crests. The central body length of *Spiniferites ristingensis*, 39 (43.0) 49 µm, is slightly smaller than that of *Spiniferites delicatus*, as given by Reid (1974).

**Remarks.** The holotype from Reid (1974) is shown in Plate 1. Additional specimens were observed in topotype material from the estuaries around the Irish Sea (Conwy Marina, Dee, Caernarvon). There was overall agreement during the Second *Spiniferites* Workshop in Ostend in 2015 that the so-called "skeletal rods", first described by Reid (1974), do not exist and are an optical illusion created by the attachment of membranes along the processes (Mertens et al., this volume).

**Occurrence**. From the lower Miocene onwards (Londeix et al. this volume).

*Spiniferites elongatus* Reid 1974

Plate 2, Figures 1–12 (holotype). Plate 5, Figures 1–6.

**Synonymy.** *Spiniferites ellipsoideus* Matsuoka 1983, see Mertens et al., this volume.

**Dimensions**. Reid (1974) – holotype: 30 × 49 µm; height antapical process: 13 µm; height apical process: 6 µm; height lateral process: 9 µm. Range: test 26 × 40 µm to 42 × 59 µm; height antapical processes: 12–16 µm; height apical processes: 6–12 µm; height lateral processes: 5–9 µm. Number of specimens measured: 15. Our measurements on topotype material from Dee Estuary – length 32 (38) 44 µm; width 20 (25) 30 µm. Number of specimens measured: 2.

**Original description (Reid 1974, p. 603).** “In polar view the test is circular. It has a thin, 0.8–1 µ wall with a smooth surface and no evidence of an apical boss. A short complex process is found at the head of the sulcus. At the junction of 1´´´´ and 3´´´ high membranous septae join two stout simple processes and at the junctions of 5´´´, 4´´´, 1´´´´ and 1P, 2´´´, 1´´´´ two high, hollow, trumpet shaped processes are found. Their multifurcate tips appear to be supported by strengthening rods. These are joined to each other and to the previous two simple processes by high septae. No sulcal plates were seen. Tabulation appears to be typical for the genus except for plate 1´´´ which is either absent or is covered by the wide septate of 2´´´ and 1P at the junction with the sulcus. Plates 1´ and 4´ are not fused but are separated by a low septa. Plate 6´´ is triangular, long and narrow. Archeopyle precingular 3´´ and reduced.”

**Additional observations.** The cysts have an elongate or ellipsoidal ambitus, and are circular in polar view. The surface is smooth to finely microgranular and no apical boss was observed. Wide flaring sutural crests are attached towards the center of the plates and leave an oval impression in the center of each plate (Plate 2, Figure 8; Plate 5, Figure 1). At the antapex, the sutural crests are high (4 µm, see Plate 5, Figure 5) and connect complex processes. The sutural crests at the apex are high as well, but lower than on the antapex (Plate 5, Figure 2), while elsewhere on the cyst, the crests are lowest, and connect simple processes. Girdle displaced by less than its own width.

**Comparison.** All elongate specimens recorded in the newly processed samples are identified as *Spiniferites elongatus* and no other elongate species such as *Spiniferites frigidus* or *Rottnestia amphicavata* were recorded. The relation to these taxa as a single morphospecies is further discussed in Van Nieuwenhove et al. (this volume).

**Remarks.** New photomicrographs of the holotype are presented in Plate 2. This species was also recorded in the newly processed topotype material from Dee Estuary.

**Occurrence**. Recorded from (mid) late Miocene or maybe middle Miocene to Recent (Londeix et al., this volume).

*Spiniferites lazus* Reid 1974

Plate 6, Figures 1–5, Plate 10, Figures 4–8.

**Synonymy.** None.

**Dimensions.** Reid (1974) – holotype: test 48×33 µm, height processes 14 µm. Range: length test: 44–58 µm, width test: 31–42 µm; depth: 31–39 µm; process length 12–25 µm; girdle width: 5–8 µm. Number of specimens measured: 32. Our measurements on topotype material from the three estuaries in the Irish Sea – length 42 (48) 56 µm; width 30 (37) 44 µm. Number of specimens measured: 4.

**Original description (Reid 1974, p. 604).** “The test is ovoid in polar view with a thick wall 1–1.5 µ which is made up of two layers of equal thickness. Sutural septae increase in height towards the processes which have process tips like wide petals in plan view. Processes surrounding the apical plates, 3´ and 2´ are in the form of a crown connected by clear fenestrate sutural septae. Girdle processes may be simple or geminal. At the junction of the postcingular and antapical plates stout simple processes are found. They may be geminal along the junction of 4´´´, 3´´´? A clear geminal process with a high fenestrate flange is found along the junction of 6´´´ an [*sic*] 1´´´´. Bases of processes are hollow, but do not pass through the test wall. A clear node or boss is found at the junction of plates 3´, 2´ and 1´. Plate 1´ and 4´ appear to be combined as there is no evidence for a suture separating them. The sulcus is slightly sigmoidal, of moderate width and with no clear sulcal plates. Tabulation is typical for the genus 3´?, 0a, 6´´, 6g, 5–6´´, 1P and 1´´´´. The archeopyle is dorsal, precingular 3´´ and reduced with a distinct elongate bell shape.”

**Additional observations.** The central body is elongated, ovoid, with an asymmetrical epi- and hypocyst, sometimes with a faint apical boss (Plate 6, Figure 3). The wall is thick (1.2 µm) and the surface is microgranular to microreticulate (Plate 10, Fig. 8). The processes are exclusively gonal and have fenestrate bases (Plate 6, Figure 4). The distal ends on processes of *Spiniferites lazus* are typically without fenestrations, trifurcate with elongated furcations ending in short bifurcate tips (Plate 6, Figure 5). The intergonal processes, as described by Reid (1974), were not observed in cysts from the topotype material. The girdle is displaced by four times its width. The archeopyle corresponds to plate 3´´ and is reduced.

**Remarks.** Cysts attributable to this species were recorded in the topotype material from Dee Estuary, Conwy Marina and Caernarvon.

**Comparison.** The fenestrate process bases are the most typical characteristic of *Spiniferites lazus*, although fenestrate bases have also been observed on *Spiniferites bentorii* and *Spiniferites hainanensis* Sun & Song 1992. *Spiniferites bentorii* is easily recognized by the pear-shaped ambitus of the central body, whereas *Spiniferites hainanensis* has a smaller number of fenestrations on the processes, while perforations are also present on the sutural crests. *Spiniferites hainanensis* furthermore has intergonal processes (Sun & Song 1992). *Spiniferites septentrionalis* has large fenestrations in the distal ends of its processes.

**Occurrence.** Middle Miocene (?) to Recent (Londeix et al. this volume).

*Spiniferites ramosus* (Ehrenberg 1837b) Mantell 1854 sensu Rochon et al. 1999

Plate 7, Figures 1–7.

**Synonymy.** *Spiniferites bulloideus* (Deflandre & Cookson 1955) Sarjeant 1970 sensu Reid 1974, Plate 2, figures 17–19. *Spiniferites ramosus* (Ehrenberg 1838) Mantell 1854 sensu Harland 1977, Plate 1, Figures 5, 6; Rochon et al. 1999, Plate 9, Figures 4–6. non *Spiniferites ramosus* var. *ramosus* sensu Davey & Rogers 1975 (Plate 1, Fig. 5).

**Original description (Reid 1974, p. 600).** “The test is oval in equatorial view, and circular to semi-circular in other views. The epitract narrowing towards the apex frequently has a shoulder at the junction of the precingular and apical plates. A thin 1µ, two layered wall with a smooth surface is ornamented with low sutural septae forming gonal processes. The processes are simple, relatively long with trifurcations that may be up to half the length of the process and have clear bifid tips. Combined geminal processes delimit the boundaries of the girdle fields and a complicated process may be found at the head of the sulcus. The junction of plates 4´´´ and 1´´´´ is marked by a germinal process which may at times be strongly developed and have two columnar stems expanding into a ‘trumpet’ shape at the distal tips. If suitably orientated the apex shows an apical horn. Girdle spiral, displaced by its own width. The girdle appears relatively wider in the smaller forms of this species. Sulcus relatively wide, showing signs of platelets. Plate 1´, 4´ appear to be fused into one plat [*sic*]. Plate 1´´´ is relatively short in length and wide laterally. Plate 6´´ is a narrow elongate triangle. Tabulation as for the genus. Archeopyle dorsal precingular 3´´.

**Additional observations.** The specimens we observed conform to those described by Reid (1974) in having exclusively gonal processes, smooth walls and low sutural crests.

**Remarks.** See Mertens et al. (this volume) for the taxonomic history.

*Spiniferites membranaceus* (Rossignol 1964) Sarjeant 1970

Plate 8, Figures 1–6.

**Synonymy.** *Hystrichosphaera furcata* var. *membranacea* Rossignol 1964; *Hystrichosphaera ramosa* var. *membranacea* (Rossignol 1964) Davey & Williams 1966; *Hystrichosphaera membranacea* (Rossignol 1964) Wall 1967.

**Dimensions.** Reid (1974) – Test 34×34 to 44×43 µm. Height of the antapical flange 2–21 µm. Process height 12–17 µm. Girdle width 5–8 µm. Number of specimens measured: 16. Our measurements – length 36 (41.4) 44 µm, width 34.4 (38.1) 42 µm, antapical flange height 12 (14) 16 µm, process height 12 (12.3) 12.8 µm. Number of specimens measured: 5.

**Original description (Reid 1974, p. 605).** “The test is circular to ovoid and slightly elongate in a posterior-anterior direction, with an apex that is marked by a clear boss. The girdle is inclined and displaced by twice its width and the sulcus is slightly sigmoid, moderately wide and consists of plates AS, RA, RS, LS and PS. The tabulation and plate pattern is typical for the genus with four apical plates and a dorsal, reduced, precingular archeopyle. The wall is two layered, each layer of equal thickness, with a microgranular to micropunctate surface ornament. Two high processes are found at the angular junctions of plates 1´´´´ and 4´´´. They are joined by an equally high membranous flange along the junction of these two plates. A sutural geminal process is found between plates 6´´´ and 1´´´´. Junctions between cingular plates are marked by similar narrow geminal processes. Plate 1P is surrounded by a more complex crown of processes joined by high septae. Apical plates are delimited from the precingulars by a crown of processes connected by membranous sutural septae. Other processes are gonal and simple. A plan view of the process tips shows narrow trifurcations from a central axis with Y-shaped terminations.”

**Additional observations.** Processes are exclusively gonal. Specimens with broken or reduced processes were often observed, lacking furcations on the distal ends (Plate 8, Figs. 1–2). The base of the sutural crests is typically observed as an undulating double line (Plate 8, Fig. 1). The cyst wall and processes are granular. The cingular displacement of two times its own width is illustrated in Plate 8, Fig. 2, which also shows a view of the slightly sigmoid sulcal area.

**Remarks.** Wall et al. (1977) remarked that *Spiniferites membranaceus* sensu Reid (1974) is probably not conspecific with that of Rossignol (1964), an observation repeated by Harland (1983). Having not been able to observe specimens from Rossignol (1964), we were not able to confirm this observation.

*Spiniferites mirabilis* (Rossignol 1964) Sarjeant 1970

Plate 8, Figures 7–9.

**Synonymy.** *Hystrichosphaera mirabilis* Rossignol 1964 [the species was first invalidly described by Rossignol (1962, p. 132, without any images), because no holotype was designated].

*Spiniferites splendidus* Harland 1979, p. 537, plate 3, figures 1–2 by Limoges et al. (this volume).

**Dimensions.** Reid (1974) – Test 44×48 to 58×60 µ. Maximum process height 15–21 µ. Girdle width 6–8 µ. Number of specimens measured: 16. Our measurements – length 40 (47.6) 60 µm, width 36 (40) 46 µm, process length 10.4 (13.9) 18 µm. Number of specimens measured: 5.

**Original description (Reid 1974, p. 606).** “Oval, slightly elongated cysts that are circular in polar view. The antapical area is ornamented by processes that are connected by a high sutural flange along the junction of plates 1´´´´ and 4´´´. At the junction of these two plates with 3´´´ and 5´´´ large processes emerge from the flange. Stout, rigid hollow gonal and sutural processes ornament the remainder of the test. The wall is thin, two layered and frequently ruptured or folded with a distinct microgranular surface. The sinistral girdle is displaced by three times its width ventrally, and the sulcus is relatively wide and shows the following plates clearly, PS, IS, LS, RS? Sutures between the right accessory and anterior sulcal plates were not seen. Tabulation is typical for the genus with a long, narrow plate 1´´´ which is delimited by low granular connections between the processes. The archeopyle is dorsal, precingular 3´´, quadrangular with rounded corners.”

**Additional observations.** Our specimens conform to the specimens described by Reid (1974).

**Remarks.** This species was observed in the Conwy Marina, Dee and Caenarvon estaries.

*Achomosphaera ramosasimilis* (Yun 1981) Londeix et al. 1999

Plate 8, Figures 10–12.

**Synonymy.** *Achomosphaera ramulifera* subsp. *ramosasimilis* Yun Hyesu 1981. “*Spiniferites ramuliferus*” sensu Reid 1974 according to Londeix et al. (this volume)

**Dimensions.** Reid (1974) – Range: 38×50 to 33×42 µ; Process length, Apical 12–17 µ, Cingular 17–25 µ, Antapical 17–25 µ. Number of specimens measured: 15. Our measurements – length 43.0 (43.8) 44.9, width 34.6 (34.9) 37.4, process length 13.3 (14.9) 16.7, wall thickness 0.7 (1.1) 1.6. Number specimens measured: 3.

**Original description (Reid 1974, p. 609).** “Recent specimens of this species have a distinct rhomboidal shape in equatorial view and ellipsoidal shape in other orientations, agreeing with Deflandre’s description of the type material. The test wall is microgranular, two layered and 1 µ thick. Hollow, wide based, sturdy, gonal spines ornament the surface reflecting a tabulation of 3–4´, 6´´, 6C, 5´´´, 0–1P´´´´. At the junction of plates 1´´´´ and 4´´´ two large processes are joined together at their bases to form a low flange. Cingular processes may also be joined together at their bases but are slender with a much shorter base. The processes are trifurcate at their distal end with the trifurcations parallel to the test wall. The processes may divide into two, each division then trifurcating to the slender extreme tips, which have a short Y-shaped bifurcation. The apex has a distinct process with two short extensions projecting from the distal end as well as the bifurcating tips. The processes appear to extend through the endophragm but are not open to the interior. Their base is covered by a thin membrane. A precingular apical-antapically elongated archeopyle is formed by the loss of plate 3´´.”

**Additional observations.** Our specimens conform to the specimens described by Reid (1974). The specimens are close to the description of *Spiniferites ramosus* sensu Rochon et al. (1999), but have no sutural traces, and narrower process bases.

**Remarks.** Reid (1974, p. 609) did not accept the genus *Achomosphaera* and considered his specimens to belong to the genus *Spiniferites*, although he mentioned the absence of sutural traces. Here, however, we assign these specimens to the genus *Achomosphaera.* The species differs from *Achomosphaera andalousiense* and *Spiniferites septentrionalis* in having no fenestrate distal ends on the processes.

**Occurrence.** We did not observe this species in the topotype material from the British estuaries. However, we encountered specimens that conform to Reid’s (1974) description of *Spiniferites ramuliferus* in type material of *Spiniferites coniconcavus* from the Pliocene Verrebroek section of Belgium (De Schepper et al. 2004). Since Reid (1974) did not depict specimens with cell content we cannot evaluate whether these specimens are reworked, and, as such, whether or not this genus is extant.

**4. Cyst wall composition**

Three *Spiniferites* species originally described in Reid 1974 (*Spiniferites belerius*, *Spiniferites elongatus* and *Spiniferites delicatus*) were measured for their cyst wall composition after isolation from Dee Estuary sediments (Figure 2). Additional species recorded by Reid in the Dee estuary sediments were also analyzed, including *Spiniferites membranaceus*, *Spiniferites* ?*ramosus* and *Spiniferites*  *ramosus*, and compared with the only published spectrum of a *Spiniferites* species (*Spiniferites pachydermus* from the Benguela upwelling region; Bogus et al. 2014) (Figure 2)*.* Five other morphologically unclassified species of *Spiniferites* were recorded and analyzed (Figure 3). In all cases, a single specimen was analyzed. As such, intraspecific variability was not assessed. However, it is expected that specimens of the same species will show similar spectra if they come from the same locality, as was demonstrated in previous studies (e.g. Mertens et al., 2015a).

All of the cyst walls are composed of a complex biopolymer, likely carbohydrate-based, that is broadly consistent with other published spectra from the same order (Gonyaulacales) (Bogus et al. 2014, Mertens et al. 2015b). Due to the complexity of the biopolymer it was not possible to definitively assign each of the IR bands. However, the differences in the intensity and position of bands in the cyst walls of the different *Spiniferites* species reflect more general variability in dinosporin composition (Figure 4).

 For all specimens (Figures 2 and 3), the region between 3600–3000 cm-1 showed a broad absorption with a maximum ~3250 cm-1 (OH stretch). Absorptions in the 3000–2775 cm-1 region (CH stretching; IR band region I) were characterized by two IR bands at 2925 cm-1 and 2860 cm-1. These absorptions are present in all specimens and reflect the contribution of CH2 and CH3 groups. The related absorptions at 1420 cm-1 and 1370 cm-1 (CH bending; IR band region III) are also present in all spectra. Additional weaker absorptions in IR band region III (e.g., 1320 cm-1 and 1280 cm-1) likely reflect C-OH deformation and C-O stretching. Absorptions between 1850–1500 cm-1 (IR band region II) are characteristic of C=C (1600 cm-1) and C=O (1705 cm-1 and 1640 cm-1) stretching, and are variable in both their presence and strength between the specimens. For example, the absorption at 1705 cm-1 is absent in the spectrum of *Spiniferites belerius*, but this IR band region contains the strongest absorption (1600 cm-1) in the entire spectrum of *Spiniferites delicatus* (Fig. 2), which influences its position in Figure 4. The absorptions between 1185–1030 cm-1 (IR band region IV) represent C-O stretching and the deformation vibrations of sugar rings at 1160 cm-1 (C-O-C asymmetric vibration), 1110 cm-1 (glucose ring stretching), 1080, 1060, 1040 and/or 1030 cm-1 (C-O stretching), as well as skeletal ring vibrations and the anomeric region (900–650 cm-1). The series of absorptions in IR band region IV provide a strong argument that a carbohydrate, probably with a β-glycosidic linkage (Pandey 1999; Kačuráková & Wilson 2001), forms the backbone of *Spiniferites* dinosporins, consistent with previous data for species from the order Gonyaulacales (Versteegh et al. 2012; Bogus et al. 2014; Mertens et al. 2015b).

 However, there is variability in IR band region IV, particularly with regards to the position of the strongest peak. Only *Spiniferites pachydermus* exhibited the absorption pattern that most closely resembles cellulose (Pandey 1999); none of the Dee Estuary *Spiniferites* specimens had their strongest absorption at 1030 cm-1 (Figure 2). The majority of the specimens that could be speciated (*Spiniferites elongatus*, *Spiniferites membranaceus*, *Spiniferites ramosus*, and *Spiniferites delicatus*) exhibited the strongest peak at 1040 cm-1 and a second peak at 1110 cm-1. The cyst wall of *Spiniferites belerius* is the most unique in region IV in that the strongest peak is centered at 985 cm-1, but with subsequent peaks at 1040, 1110 and 1160 cm-1, similar to *Spiniferites pachydermus*; it is the only specimen that contains such a shifted primary absorption, and an overall broader series of absorptions, which is reflected in its position in Figure 4. The only other specimen with a dominant absorption in region IV not centered at 1040 cm-1 is *Spiniferites* ?*ramosus*, where it is instead centered at 1080 cm-1. This is more similar to many of the unspeciated *Spiniferites* specimens (i.e., *Spiniferites*  spp. 3, 4, and 5; Fig. 3) that also exhibit their strongest absorption at 1080 cm-1. *Spiniferites* spp. 1 and 2 exhibit the dominant absorption at 1040 cm-1, which is more in line with the speciated specimens.

 All species measured from the topotype material have a more pronounced peak in the range of 1600-1705 cm-1 (region II) compared with the spectrum of *Spiniferites pachydermus* from the Benguela upwelling region. This is probably due to environmental differences between the surface waters of the two regions. It has been shown for other species that different water mass conditions lead to different cyst wall chemistries (Mertens et al., 2015a). An analytical factor cannot be fully ruled out, however, although limited published data using a chitin standard (described in Bogus et al., 2014) comparing spectra produced in the respective laboratories demonstrated that the spectra were comparable.

 It is clear from both the morphological examination and chemical analysis that *Spiniferites delicatus*, *Spiniferites belerius*, and *Spiniferites membranaceus* are quite distinct (Plates 3, 4 and 8, Fig. 4). The chemical composition of *Spiniferites elongatus* suggests that it is most similar to *Spiniferites membranaceus*, and that *Spiniferites ramosus* and *Spiniferites* ?*ramosus* are most similar to each other (Fig. 4). In conjunction with the morphological determination for *Spiniferites ramosus* and *Spiniferites* ?*ramosus*, it is possible that they are, in fact, the same species (*Spiniferites ramosus*); unfortunately, it is not currently possible to be conclusive regarding this.

 If the visually defined species’ cyst wall chemistries are used as a guide, there is some indication that dinosporin composition can constrain the unspeciated *Spiniferites* specimens. This is particularly evident for *S.* spp. 2, as it plots very close to *Spiniferites ramosus* and close to *Spiniferites* ?*ramosus*, suggesting that this specimen may not be a different species, and that the difficulty of its morphological classification is caused by the wide range in morphological variability. However, the other unspeciated *Spiniferites* specimens seem to cluster nearer to each other with *Spiniferites*  spp. 4 and 1 and *Spiniferites*  spp. 3 and 5 showing greater proximity (Fig. 4). This could indicate that each of these couples is a different species, or that all four represent morphological variations of the same species. It is currently not possible to determine this, but it is clear that their dinosporins are different from the speciated *Spiniferites*. The cyst wall chemistry also demonstrates the ambiguity in defining species of this genus.

**5. Conclusions**

The *Spiniferites* species described by Reid (1974), *Spiniferites belerius*, *Spiniferites delicatus*, *Spiniferites elongatus* and *Spiniferites lazus*, were re-studied to improve their morphological descriptions and iconography. The holotypes of *Spiniferites delicatus* and *Spiniferites elongatus* were re-photographed to improve the image resolution. Both of the latter species, together with *Spiniferites belerius* and *Spiniferites lazus*, were also recorded in newly processed topotype material from three British estuaries (Dee, Conwy Marina and Caernarvon). *Spiniferites belerius* is characterized by its size and the trumpet-shaped process, and its position has been assessed as the junction of plates 1´´´´ and 1p. *Spiniferites delicatus* was well-defined in the original description but the distinction with *Spiniferites ristingensis* needs further investigation. Specimens of *Spiniferites elongatus* and *Spiniferites lazus* posed no problems with the original descriptions.

Specimens of *Spiniferites membranaceus*, *Spiniferites mirabilis*, *Spiniferites ramosus* (called *S. bulloideus* by Reid 1974) were also identified in and illustrated from the topotype material from the British estuaries. The species *Spiniferites hyperacanthus* and *Spiniferites pachydermus* were not recorded. *Spiniferites ramuliferus*, now known to be *Achomosphaera ramosasimilis* was not recorded in the topotype samples, but cysts corresponding to the description of this species as given by Reid (1974) were identified in material from the Pliocene of Belgium.

 Along with the taxonomic re-investigation, the cyst wall composition of *Spiniferites* specimens isolated from Dee Estuary sediments, both visually speciated (*Spiniferites belerius*, *Spiniferites ramosus*, *Spiniferites* ?*ramosus*, *Spiniferites delicatus*, *Spiniferites elongatus*, *Spiniferites membranaceus*) and not (*Spiniferites* spp. 1–5), were compared. The results showed variations in the *Spiniferites* species’ dinosporin compositions, supporting the suggestion that cyst wall chemistry may be species specific and evident when other factors are controlled (i.e. geography and environmental conditions). The cyst wall chemistries of *Spiniferites delicatus*, *Spiniferites belerius*, and *Spiniferites membranaceus* were the most distinct, suggesting that the morphological assignation for the different species is supported by the cyst wall chemistry. It also provides some impetus for clarifying the species’ descriptions for *Spiniferites belerius* and *Spiniferites delicatus* in particular.

 The unspeciated *Spiniferites* specimens were mostly difficult to constrain chemically, as they did not show clear overlap with the speciated spectra. Overall, the visually unidentifiable specimens represent examples of the ambiguity in speciating *Spiniferites* specimens morphologically and chemically. However, *Spiniferites* spp. 1, and 3 to 5 clustered nearer to each other in terms of cyst wall chemistry, suggesting that they may be the same species, but with enough morphological variation to make their visual identification difficult. Furthermore, *Spiniferites* spp. 2 plotted very close to *Spiniferites ramosus*, indicating that this specimen may, in fact, be the same species.

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**Author contributions**

PG, KM, KB, FM & SL conceived the study; PG studied and photographed the holotypes in Sheffield; FM provided the sediments from topotype localities; PG processed the new sediment samples; KM & PG studied and photographed the cysts from topotype localities with light microscope; KM & PG processed the residues, picked cysts for FTIR; PG, KM & HV did spectral analysis for cyst wall composition; KB interpreted the spectra and the relation to dinocyst wall chemistry; PG, KM, KB & SL wrote the paper and PG, KM & SL made the plates.

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**Plates**

Plate 1. Figures 1–16: *Spiniferites delicatus*. Holotype 12 ABSL 1290.192(5). 1–7: High focus, incrementally lowering; 8–9: Optical sections; 10–16: Low focus, incrementally lowering. Scale bar = 10 µm.

Plate 2: Fig. 1–12: *Spiniferites elongatus*. Holotype 140K3 1210.352(12). 1–3: High focus, incrementally lowering; 4-5: Optical sections; 6–12: Low focus, incrementally lowering. Scale bar = 10 µm.

Plate 3: 1–12: *Spiniferites belerius*. 1–3: Dee Estuary slide 1/3 sp. 1, incrementally lowering focus. 4: Dee Estuary slide 1/3 sp. 2, high focus. 5–6: Dee Estuary slide 3/3; 5: optical section, 6: low focus. 7–10: Dee Estuary slide B/B sp. 1, incrementally lowering focus; 10: optical section. 11: Dee Estuary slide B/B sp. 2, high focus. 12: Dee Estuary l.l, detail of wall structure. Scale bar = 10 µm.

Plate 4: Fig. 1–9: *Spiniferites delicatus*. 1–6: Celtic Sea 9-8 St. 8 5100; 1–2 High focus, incrementally lowering; 3: Optical section; 4–6: Low focus, incrementally lowering. 7–9: Celtic Sea St. 8 11 1 99 0 cm cs 5–6; 7: High focus; 8: Optical section; 9: Low focus. Scale bar = 10 µm.

Plate 5: Fig. 1–6: *Spiniferites elongatus*. Dee estuary. 1: High focus; 2: Optical section; 3–6: Low focus, incrementally lowering. Scale bar = 10 µm.

Plate 6: Fig. 1–5: *Spiniferites lazus*. 1–3: Celtic Sea St. 5 4100 cs 9–4, sp. 9; 1–2: High focus, incrementally lowering; 3: Optical section. 4: Celtic Sea St. 5 4100 cs 9–4, sp. 10, detail of perforations at process base; 5: Celtic Sea St. 5 4100 cs 9–4, sp. 2, detail of process terminations. Scale bar = 10 µm.

Plate 7. 1–6: *Spiniferites ramosus*. 1–3: Dee Estuary 3/3; incrementally lowering focus. 4–6: Celtic Sea St. 5 4100 cs 9–4; 5: high focus; 6: optical section; 7: low focus. Scale bar = 10 µm.

Plate 8. 1–6: *Spiniferites membranaceus*. 1–2. Dee Estuary, SEM-images. 3–6: Celtic Sea St. 6 3100 cs 8–4, 3–4, high focus, 5 optical section, 6 low focus. 7–9: *Spiniferites mirabilis*. Celtic Sea St. 6 4100 cs 9–3. 7: high focus; 8: optical section; 9: low focus. 10–12: *Achomosphaera ramosasimilis*. Verrebroek dock section. Incrementally lowering focus. Scale bar = 10 µm.

Plate 9. 1–8: *Spiniferites ristingensis*. Britanny, Vilaine Estuary. 1: detail of external wall texture; 2: axial detail of process termination; 3: detail of wall at the archeopyle margin; 4: detail of cingular processes; 5: oblique view of process termination; 6: antapical view; 7: dorsal view; 8: lateral view. Scale bar = 10 µm.

Plate 10. 1–3: *Spiniferites coniconcavus*. Pliocene Verrebroek section, Belgium. 1: antapical view; 2: dorsal view on archeopyle; 3: oblique dorsal view on archeopyle. 4–8: *Spiniferites lazus*. 4: right lateral view; 5: dorsal view with archeopyle visible; 6: left lateral view; 7: antapical view; 8: detail of wall surface. Scale bar = 10 µm, except in 8, where it is 2 µm.

**Figure captions**

Figure 1. Location of the samples studied by Reid (1974) (red dots). Topotype localities from the Irish Sea are indicated as 1 (Dee estuary), 2 (Conwy Marina) and 3 (Caernarvon).

Figure 2. FTIR spectra of morphologically identifiable *Spiniferites* species from Dee estuary sediments. The spectrum colors correspond to Figure 3. IR band regions used in the relative strength comparison are (I) 3010–2775 cm-1, (II) 1850–1500 cm-1, (III) 1500–1185 cm-1, and (IV) 1185–860 cm-1. The published spectrum of *Spiniferites pachydermus* (core GeoB4804, Benguela upwelling region) is shown for comparison (Bogus et al. 2014).

Figure 3. FTIR spectra of morphologically unidentifiable species of the genus *Spiniferites* (*Spiniferites* spp.) from Dee estuary sediments. The spectrum colors correspond to Figure 3. IR band regions used in the relative strength comparison are (I) 3010–2775 cm-1, (II) 1850–1500 cm-1, (III) 1500–1185 cm-1, and (IV) 1185–860 cm-1.

Figure 4. Relative IR band comparison between morphologically identifiable *Spiniferites* species, and morphologically unidentifiable cysts of the genus *Spiniferites*. The IR regions correspond to (I) 3010–2775 cm-1, (II) 1850–1500 cm-1, (III) 1500–1185 cm-1, and (IV) 1185–860 cm-1.

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