

The trace fossil *Polykampton recurvum* n. isp. (sequestrichnia) from the Maastrichtian–Paleocene deep-sea deposits of NW Italy

Alfred Uchman^{1*}  and Bruno Rattazzi²

¹Jagiellonian University, Institute of Geological Sciences, Gronostajowa 3a; 30-387 Kraków, Poland <alfred.uchman@uj.edu.pl>

²Museo Paleontologico di Crocefieschi, Via alla Chiesa 12, 16010 Crocefieschi (Genova), Italy <museocrocefieschi@libero.it>

Abstract.—*Polykampton recurvum* n. isp. is the sixth ichnospecies of the ichnogenus *Polykampton* Ooster, 1869. It is a horizontal structure composed of a median cylindrical tunnel and narrow, usually back-curved lateral lobes located in alternating position. It occurs 2–3 cm below the top of single beds in the Maastrichtian–Paleocene deep-sea turbiditic marlstones of the Monte Antola Unit in the Northern Apennines. The lobes of *P. recurvum* n. isp. are actively filled with gray mudstone from above through the permanently open median tunnel. The trace fossil belongs to the category sequestrichnia, which is typical of oxygenated deep-sea environments characterized by seasonal or episodic supply of organic matter into a generally oligotrophic environment. *P. recurvum* n. isp. was produced by a “worm,” probably a polychaete, which adapted to seasonal or only episodic supply of organic matter to the deep-sea floor. The tracemaker stored the organic-rich mud in the lobes for nutrition during times of low organic matter availability on the seafloor.

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Introduction

The trace fossil *Polykampton* Ooster, 1869, typified by *P. alpinum* Ooster, 1869 from the Paleogene flysch deposits of Switzerland, was almost forgotten for more than a century and treated as a monospecific ichnogenus (Häntzschel, 1975; Seilacher, 2007). Discoveries of four new ichnospecies of *Polykampton* from Cretaceous–Oligocene flysch deposits (Wetzel and Uchman, 1997; Uchman and Rattazzi, 2018; Uchman et al., 2019, 2020) showed that this ichnogenus is more diverse and widespread than previously thought. In this paper, a sixth named ichnospecies of *Polykampton* is described and interpreted. The new ichnospecies derives from the Maastrichtian–Paleogene deep-sea deposits of the Northern Apennines in Italy. The objectives of this study are to add to the knowledge on: (1) the general increase in diversity of deep-sea trace fossils since the Late Cretaceous, (2) the diversification of their nutritional strategies since the Late Cretaceous (Uchman, 2004), and (3) the competitive behavior of burrowing invertebrates in generally oligotrophic environments.

Geological setting

The study area is characterized by the Monte Antola Unit of the Northern Apennines, specifically, in the Val Borbera region and its surroundings, north of Genova in northwestern Italy (Fig. 1). The Monte Antola unit is probably an allochthonous tectonic slab of Ligurid units (e.g., Marroni et al., 2001 and references

therein). The described trace fossil occurs in the stratigraphically higher part of the slab, which is composed of the Bruggi-Selvapiana Formation overlain by the Pagliaro Formation.

The Bruggi-Selvapiana Formation (late Campanian–late Maastrichtian) is a c. 500 m thick succession of thick- and very thick-bedded hybrid turbidites. Their lower part is composed of siliciclastic material, and their upper part consists mainly of marlstones (marlstone understood as hard rock of the composition as in marl) in the upper part, which alternate with subordinate thin-bedded sandstones and shales (Abbate and Sagri, 1967). In the Borbera Valley, this unit is thinner than elsewhere because it pinches out here (Levi et al., 2006).

The Pagliaro Formation (Bellinzona et al., 1971; Marroni et al., 2001, 2002; Levi et al., 2006) is about 300–400 m thick (Abbate and Sagri, 1967) and is dated to nannoplankton NP1–NP5 zones of the early Paleocene to early late Paleocene (Marroni et al., 2001) or to the CC25b–NP5 zones of the late Maastrichtian–early late Paleocene (Levi et al., 2006; Catanzariti et al., 2007). In the lower to middle part, the Pagliaro Formation consists mainly of thin- to thick-bedded turbidites, whose beds are composed of sandstones and/or siltstones becoming gradually replaced by turbiditic marlstone and/or shale. At the top of the beds, the shales represent partly the pelagic and hemipelagic background sedimentation. They are commonly dark gray and noncalcareous. The silty or muddy turbiditic marlstones are massive and light gray, dark gray, or pale rose in color. Some of these beds are up to a few meters thick. The sandstones are rich in plant detritus. Commonly, thinner beds display the Ta–d Bouma intervals. Locally, the beds thicken in upward trends in packages about 10 m thick. In the upper part of the

*Corresponding author

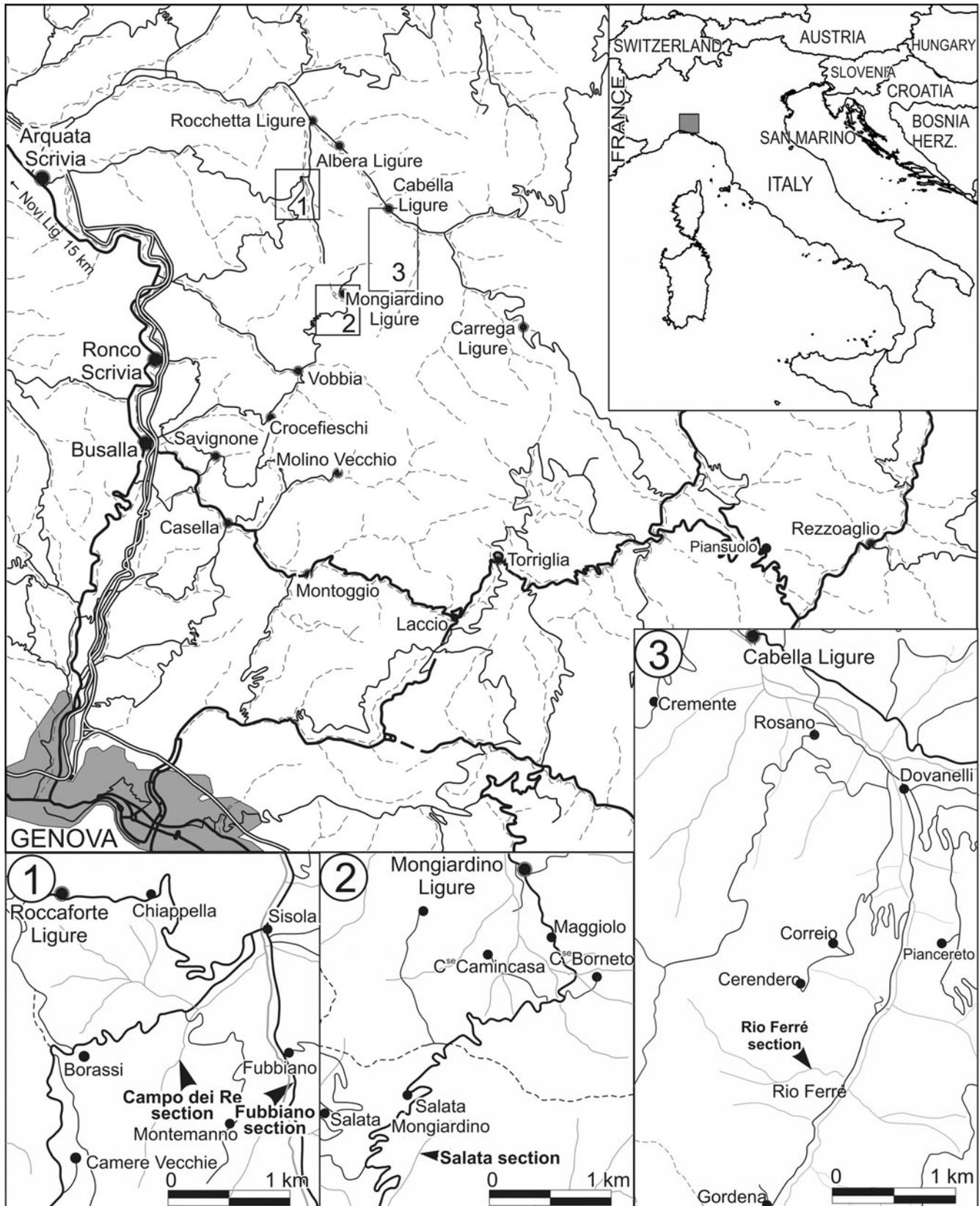


Figure 1. A general location map with inserts containing indications of the sections in detail. (1) Campo dei Re section and Fubbiano section. (2) Salata section. (3) Rio Ferré section. Location of the study region is indicated by the gray quadrangle in the upper right insert.

formation, the marlstone beds are less common. The formation is gently tectonically deformed, with folds, faults, and internal detachments, especially in the middle and upper parts (Marroni et al., 2002; Levi et al., 2006). Trace fossils of the Pagliaro Formation are very diverse and locally abundant; they are typical of the deep-sea *Nereites* ichnofacies (Uchman, 2007; Uchman and Rattazzi, 2018; Uchman et al., 2019).

The Pagliaro Formation is overlain unconformably by lower Oligocene fan-delta conglomerates of the Savignone Formation and locally by other Oligocene sediments, mainly the Ranzano Formation (e.g., Gelati and Gnaccolini, 1978; Ghibauda et al., 1985; Gnaccolini, 1988). The conglomerates pass upward into the kilometer-thick series of turbidites of the Monastero Formation (Bellinzona et al., 1971; Mutti et al., 1995; Marroni et al., in press). In the lower part, the turbidites are locally conglomeratic, while pelitic facies prevail in the upper part. Locally, pebbly mudstones and slump deposits are present. The calcareous nannoplankton matches the NP23 or NP24 zones of late Rupelian to early Chattian (Marroni et al., in press). These lithostratigraphic units belong to the filling of the episutural Tertiary Piemonte Basin.

The described *Polykampton* occurs in five sections, four in the middle part of the Pagliaro Formation (Fig. 1), the Campo dei Re section (44°39.440'N, 009°02.438'E), the Salata (44°37.063'N, 009°03.042'E) and Fubbiano (44°40.086'N, 009°03.035'E) sections, and one section in the Bruggi-Selvapiana Formation in the Rio Ferré (44°38.283'N, 009°06.217'E). At the first locality, it was found in a marlstone bed outcropping in a few-meters-high escarpment on the right side of a ravine (Fig. 2). Most specimens derive from this bed. In the Salata section, turbiditic deposits a few tens of meters thick are exposed in a steep ravine. The trace fossil was found in a marlstone bed on the left bank of the stream close to the bottom of the ravine (Fig. 2). In the Fubbiano section (an escarpment in the river valley) and Rio Ferré sections (a stream bed in a ravine), specimens of the described *Polykampton* had been found occasionally in previous years. Those specimens derive from beds that have recently been covered, so detailed sections cannot be presented.

Materials and methods

Sections at Campo dei Re and Salata were measured bed by bed, and collected specimens were hand prepared. For better contrast between host rock and trace fossils, some split surfaces were oxidized with a weak acid. The same effect is obtained after at least a few months of weathering in natural conditions. The size measurements are made by means of mechanical calipers.

Repositories and institutional abbreviations.—Type material and other specimens examined in this study are deposited in the following institutions: The Crocefieschi Museum near Genova in Italy and the Nature Education Centre of the Jagiellonian University (CEP) – Museum of Geology (Kraków, Poland). Specimens of the new *Polykampton* ichnospecies labeled with four-digit numbers are housed in the Crocefieschi Museum. Specimens in the Nature Education Centre of the Jagiellonian University (CEP) – Museum of Geology are under the acronym INGUI249P. Several specimens are composed of two pieces of a split bed (labeled as “a” and “b” after the

number [in CEP] or as “bis” for one of them, which is the counterpart [in the Crocefieschi Museum]).

Systematic ichnology

Ichnogenus *Polykampton* Ooster, 1869

Type ichnospecies.—*Polykampton alpinum* Ooster, 1869 from the Paleogene flysch deposits of Switzerland, by original designation.

Diagnosis.—Horizontal, ribbon-like structure composed of a median cylindrical tunnel and complex leaf-like lobes (Uchman et al., 2020, modified from Uchman and Rattazzi, 2018 and Uchman et al., 2019).

Remarks.—The ichnogenus *Polykampton*, aside from its type ichnospecies *Polykampton alpinum*, includes *P. eseri* (Unger, 1850) (Wetzel and Uchman, 1997), *P. cabellae* Uchman and Rattazzi, 2018, *P. guberanum* Uchman, Wetzel, and Rattazzi, 2019, *P. multifiellatum* Uchman, Wetzel and Rattazzi, 2019, and *P. georgianum* Uchman et al., 2020. They are known from Upper Cretaceous to Oligocene flysch deposits, so far only in Europe and the Caucasus region.

Polykampton recurvum new ichnospecies
Figures 3–6

Holotype.—Specimen 7335 (Fig. 3.1) from the Pagliaro Formation in the Campo dei Re section.

Paratypes.—Paratype 1, INGUI149P230b (Fig. 3.2–3.4). Paratype 2, specimen 7333bis (Fig. 3.5). Paratype 3, INGUI149P234b (Fig. 3.6), all from the Pagliaro Formation in the Campo dei Re section.

Diagnosis.—Horizontal structure composed of a median cylindrical tunnel and narrow, usually back-curved, sparse, lateral lobes arranged alternately.

Description.—The median tunnel is straight, gently curved, or slightly winding, 0.7–5 mm wide (mean 2.3 mm, n = 45; see Fig. 7 for morphometric parameters). In some specimens, width of the median tunnel changes gradually toward one end by ~20–40%. The length of the median tunnel was observed for ~12–85 mm (mean 39.7 mm, n = 45). Its natural termination is unknown because the tunnel plunges into or emerges from the rock or is broken at the edge of a specimen. The width of the median tunnel is usually the same or larger than the width of the lobes. Rarely, the contrary situation is seen (Fig. 4.3), but in such cases this could be an effect of intersection if the tunnel runs at a slightly different level from the lobes. The variable morphometric parameters are shown in Figure 7.1 and the data in Table 1. Selected statistic relationships between some morphometric parameters are presented in Figure 7.3–7.8.

The lateral lobes are located alternately positioned on both sides of the tunnel. They show a constant width and a semicircular termination. The width of individual lobes may differ within the same specimen. In all specimens (n = 45), the width of the

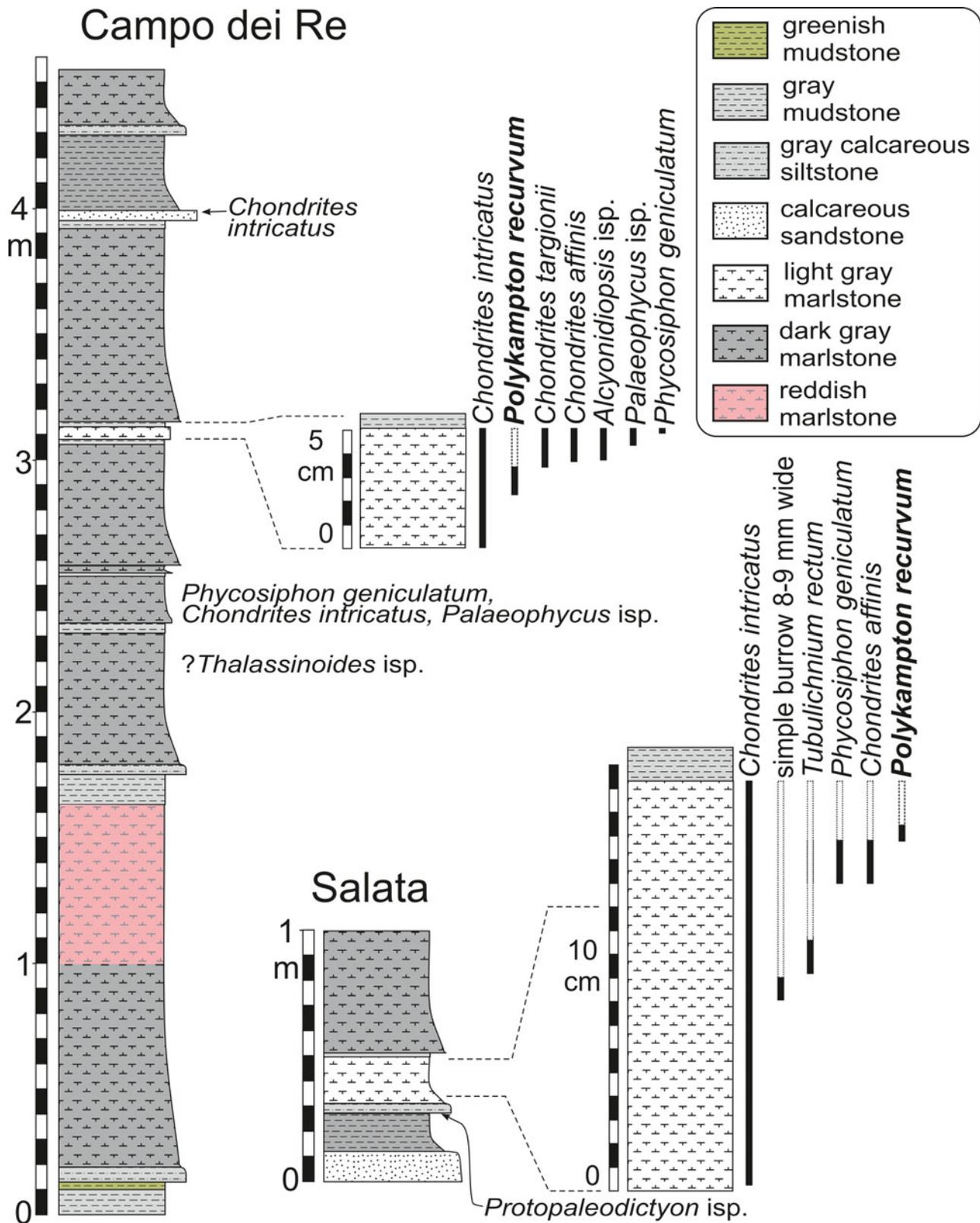


Figure 2. Fragments of sections of the Pagilaro Formation at Campo dei Re and Salata with indication of trace fossils and distribution in the bed bearing *Polykampton*.

widest lobes ranges from 1 to 4.5 mm (mean 2.2 mm), and the width of the narrowest lobes ranges from 0.7 mm to 4 mm (mean 1.8 mm). The narrower lobes are usually more frequent

in one part of the specimen than in the other (Figs. 4.2, 4.4, 5.1, 5.8, 6.2, 7.2). The length of the lobes may differ within a specimen. The longest lobes reach 3.5–18 mm (mean 5.3 mm)

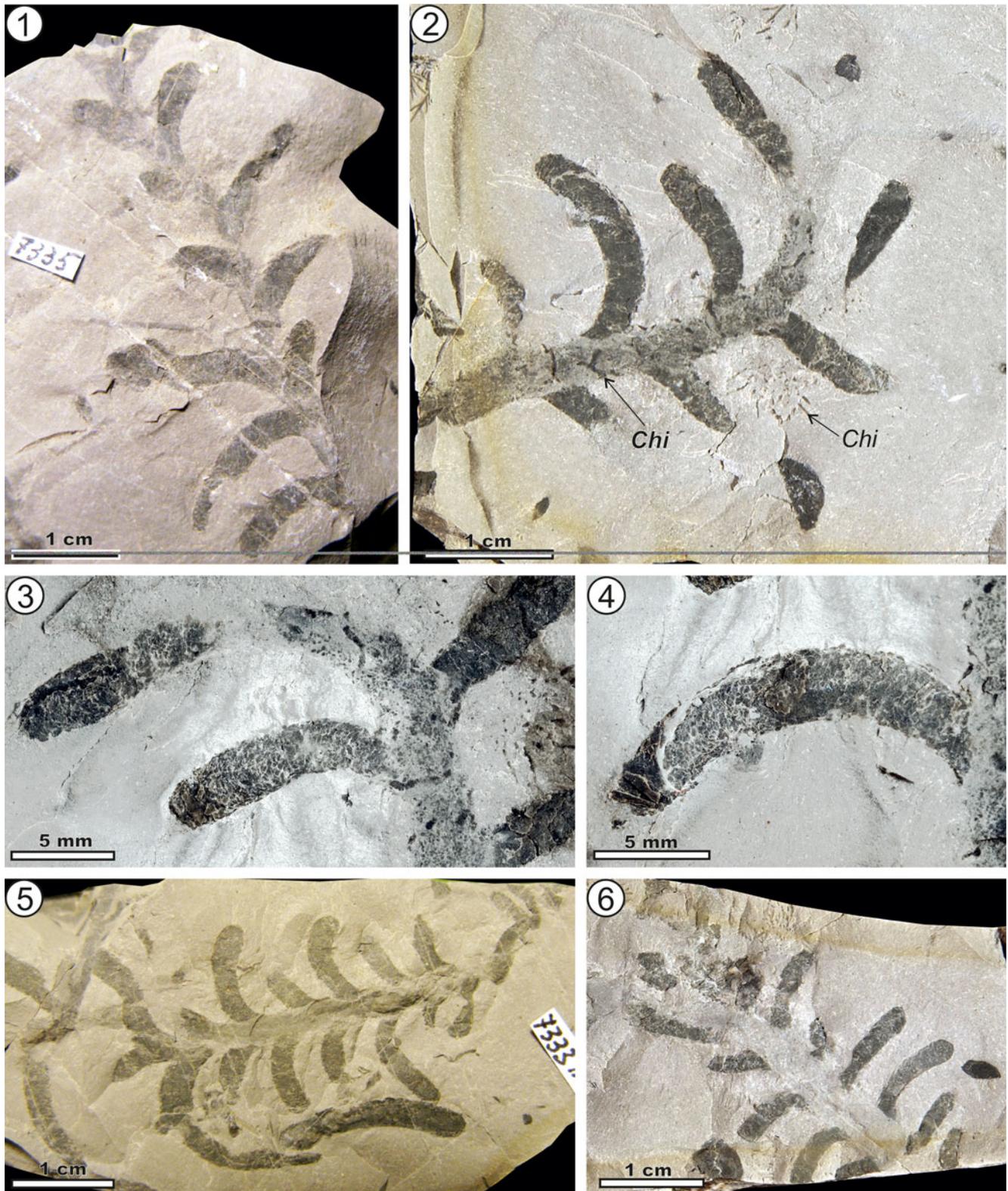


Figure 3. Types of *Polykampton recurvum* n. isp. from the Pagliaro Formation in the Campo dei Re section. (1) The holotype, specimen 7335. (2) Paratype 1, INGUJ149P230b. *Chi* = *Chondrites intricatus*. (3, 4) Details of (2). (5) Paratype 2, specimen 7333 bis. (6) Paratype 3, INGUJ149P231b.

and the shortest lobes are 1–6.5 mm long (mean 4.2 mm). The spacing between the near lobes ranges from 1 to 19 mm (mean 4.1 mm) and may vary significantly within a specimen.

The maximum width of the trace fossil (measured perpendicularly to the median tunnel) ranges from 8 to 38 mm (mean 17.8 mm, $n = 45$). The number of lobes in better-developed

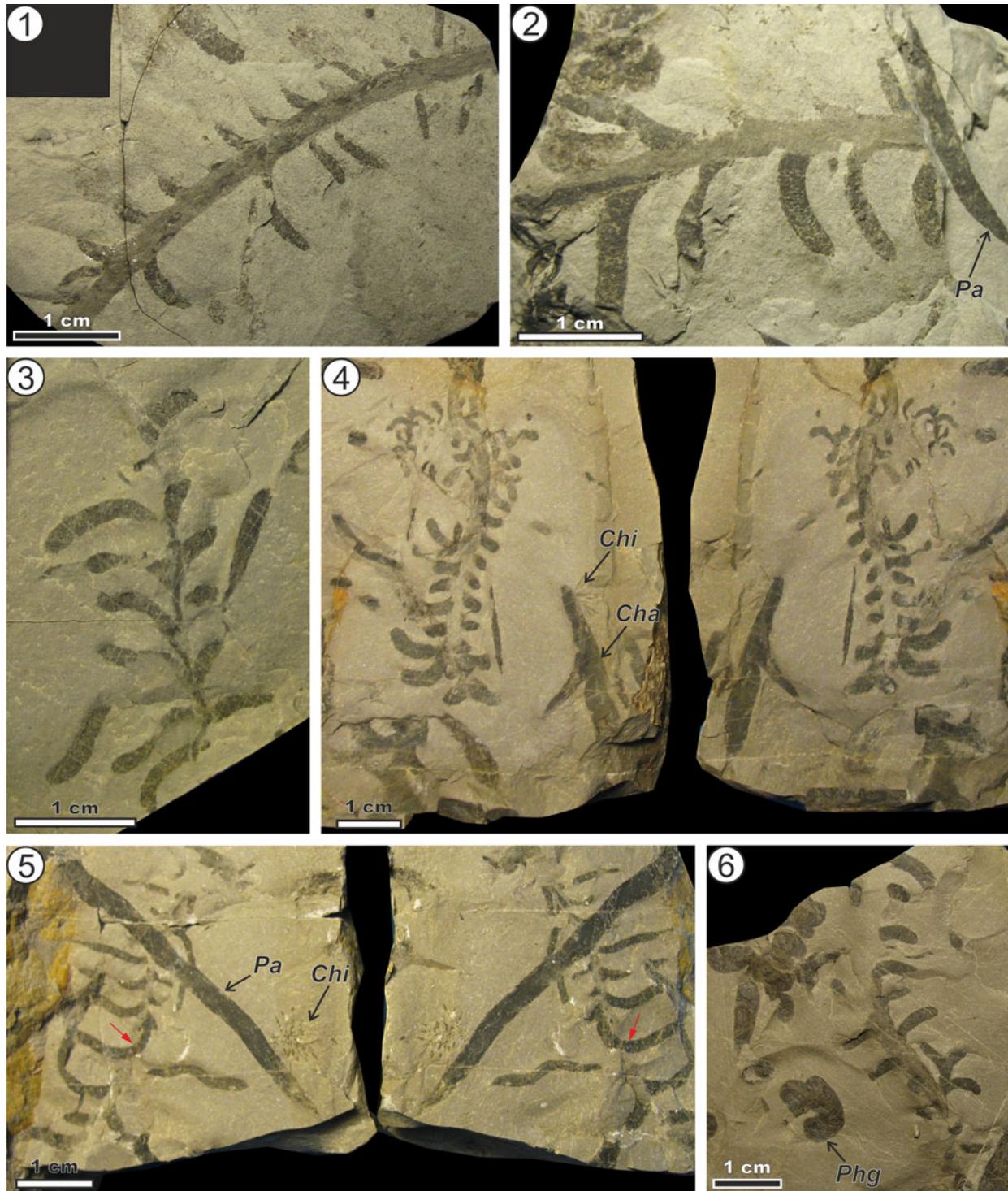


Figure 4. Selected specimens of *Polykampton recurvum* n. isp. from the Pagliaro Formation in the Fubbiano (1, 2) and Campo dei Re (3–6) sections. *Cha* = *Chondrites affinis*; *Chi* = *Chondrites intricatus*; *Pa* = *Palaeophycus* isp.; *Phg* = *Phycosiphon geniculatum*. (1) Specimen 6755. (2) 6756bis. (3) 7137. (4) 7213 and 7213 bis. (5) 7145 and 7145bis. Red arrows point to intersection of lobes with the median tunnel. (6) 7136bis.

specimens ranges from 8 to 24 (mean 13.3, $n = 24$). This gives 12–40 lobes/20 cm (mean 23.8 lobes/10 cm).

The lobes emanate tangentially from the mid or basal flanks of the median tunnel and run in the same direction at an angle of $\sim 45^\circ$ to the tunnel axis, but shortly after (within < 2 mm) they turn to a more perpendicular orientation and form an arc. The

arc may be prolonged backward (i.e., recurved). In most specimens, the lobes form regular arcs having a similar curvature, but some are slightly winding (Figs. 3.5, 4.6, 5.1, 5.4, 6.6). Rarely, the course of some lobes continues in the direction of emanation from the median tunnel (Figs. 3.2, 4.5, 6.2); that is, they are not recurved. Even more rarely, the lobes can be curved

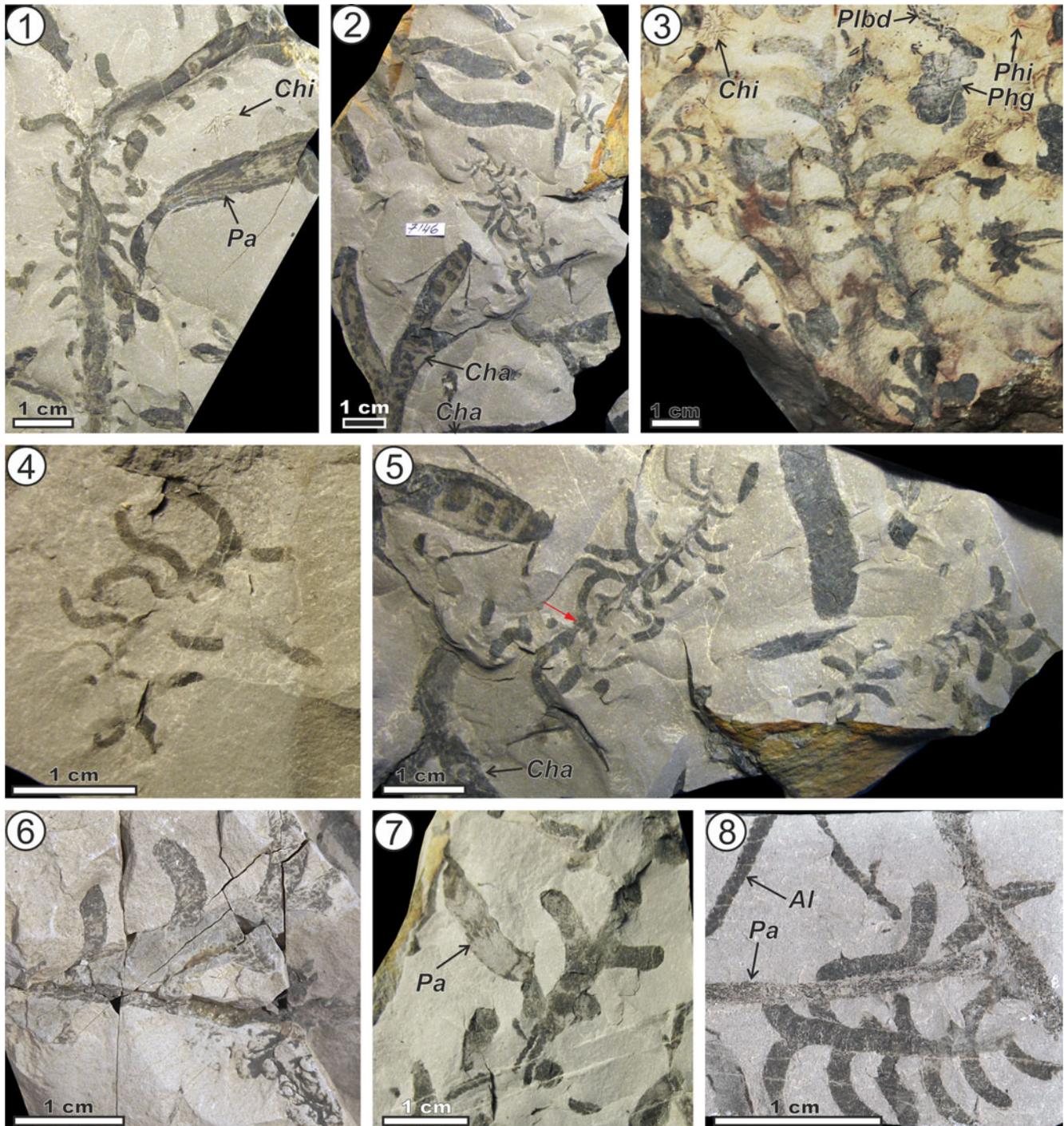


Figure 5. Selected specimens of *Polykampton recurvum* n. isp. from the Pagliaro Formation in the Campo dei Re section (1, 2, 4–8) and from the Bruggi-Selvapiana Formation in the Rio Ferré section (3). Al = *Alcyonidiopsis* isp.; Cha = *Chondrites affinis*; Chi = *Chondrites intricatus*; Pa = *Palaeophycus* isp.; Phg = *Phycosiphon geniculatum*; Phi = *Phycosiphon incertum*; Plbd = *Planolites* with filling bioturbated with *Chondrites* (“Bandchondriten”). (1) Specimen 7148bis. (2) 7146. (3) 6718. (4) 7139. (5) Detail of (2), 7146; red arrow points to intersection of lobes with the median tunnel. (6) INGUJ149P244. (7) 7151. (8) INGUJ149P235.

toward the median tunnel shortly after emanation and miss the tunnel at a slightly different level (Figs. 4.5, 5.5).

Rarely, some lobes are branched, with one or two branches, which are straight or curved back (Figs. 4.6, 6.6). Exceptionally, about half of the lobes in specimen 7146 are branched (Fig. 5.5). The lobes and the median tunnel can be on slightly different levels; that is, the lobe can run deeper than the tunnel. Therefore,

on some parting surfaces, incomplete lobes may be visible whose length can be underestimated.

The trace fossil is filled with noncalcareous gray mudstone, which contrasts with the beige color of the surrounding marlstone. Filling of the median tunnel is lighter than filling of the lobes. Not rarely, sediment in the filling of the lobes is pelleted and slightly meniscate (Figs. 3.2–3.4,

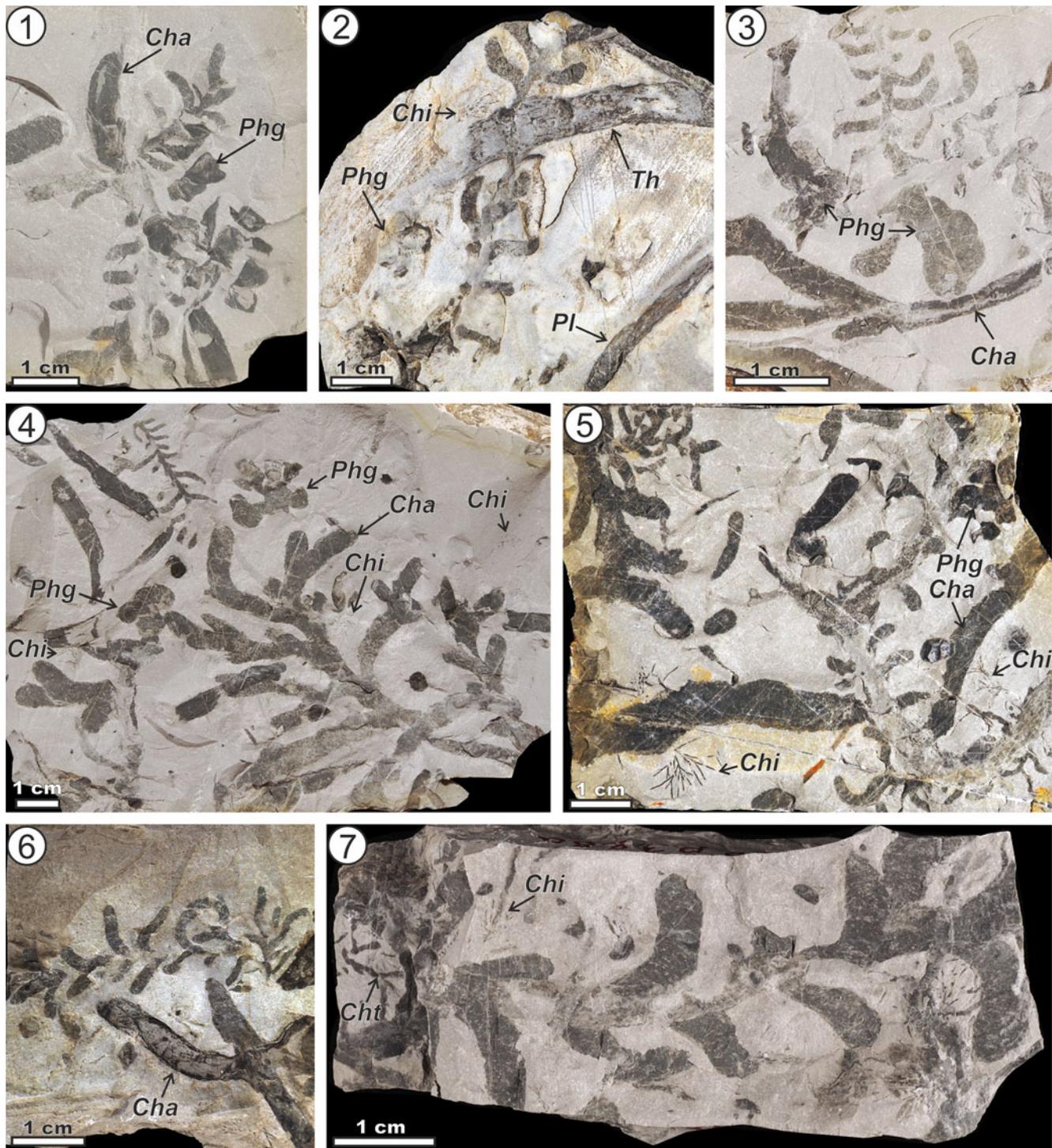


Figure 6. Selected specimens of *Polykampton recurvus* n. isp. from the Pagliaro Formation in the Campo dei Re (1–6) and the Salata (7) sections. *Cha* = *Chondrites affinis*; *Chi* = *Chondrites intricatus*; *Cht* = *Chondrites targionii*; *Phg* = *Phycosiphon geniculatum*; *Pl* = *Planolites* isp.; *Th* = *Thalassinoides* isp. (1) INGUJ149P237a. (2) INGUJ149P233. (3) INGUJ149P238. (4) INGUJ149P239a. (5) INGUJ149P231b. (6) INGUJ149P390. (7) INGUJ149P388b.

4.2, 6.7). The pellets are indistinct, elliptical in outline, less than 0.5 mm in diameter. The menisci are manifested as shallow, densely packed convex-outward arcs on the surface. Exceptionally, in one specimen (ING149P235), pelleted and meniscate sediment also fills the median tunnel (Fig. 5.8).

Etymology.—*Recurvus* (Latin) means bent back. This corresponds to the shape of the lobes.

Material.—Thirty-four specimens (6476, 7079, 7136, 7137, 7138, 7139, 7145, 7146, 7147, 7148, 7151, 7182, 7183, 7213, 7321, 7334, 7336, 7372, 7390, 7455, 7456, INGUJ149P231,

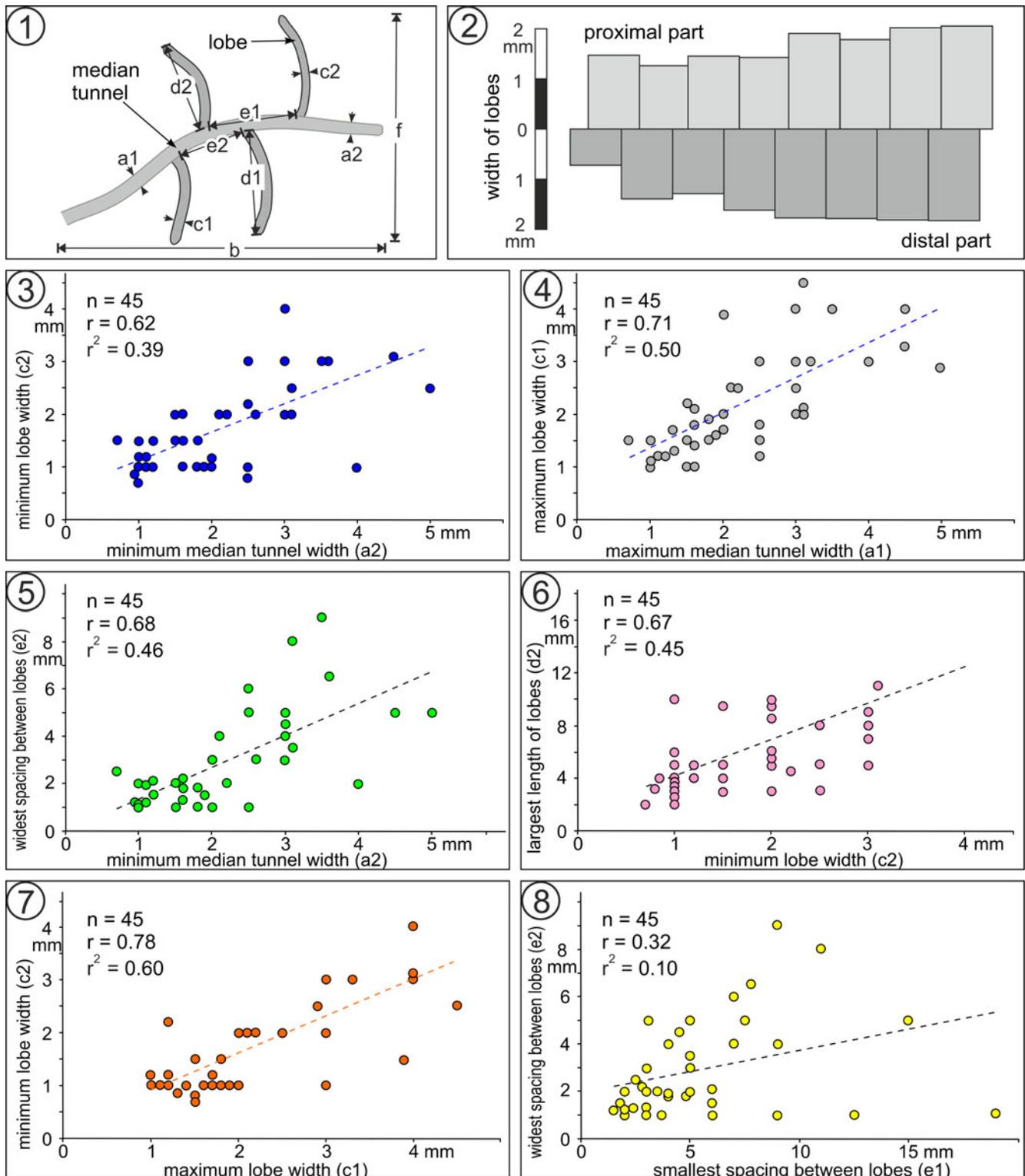


Figure 7. (1) Morphometric parameters of *Polykampton recurvum* n. sp. (2) Width of subsequent lobes on both sides in specimen 7213. (3–8) Selected statistical relationships between some morphometric parameters.

233, 235, 237, 238, 239, 243, 244, 246, 247, 387, 390, 391) from the Campo dei Re section (Pagliaro Formation), five specimens (6445, 6933, INGUI149P387, 388, 408) from the Salata

section (Pagliaro Formation), two specimens (6755, 6756) from the Fubbiano section (Pagliaro Formation), and one specimen (6718) from the Rio Ferré section (Bruggi Selvapiana Member).

Table 1. Morphometric parameters of *Polykampton recurvum* n. isp., n = 45. For the parameter definitions, see Figure 7.1.

Parameter	a1	a2	b	c1	c2	d1	d2	e1	e2	f
Range (mm)	0.7–5	0.7–4.5	11.5–83	1–4.5	0.7–4	3.5–17	2–17	1.8–5.6	1–8	8–38
Mean (mm)	2.3	2.2	39.7	2.2	1.8	9.6	6.3	5.3	2.9	17.8

Remarks.—*Polykampton recurvum* n. isp. shows all features of *Polykampton* as presented in the diagnosis of the ichnogenus, first being the median tunnel and lateral lobes as the principle ichnotaxobase. The shape and arrangement of the lobes is the ichnotaxobase for the ichnospecies of *Polykampton*. Compared with the other ichnospecies, it is most similar to *P. eseri* from the Late Cretaceous and Eocene of the Alps (Uchman, 1999; Wetzel and Uchman, 1997). However, *P. eseri*, known only from a few specimens, is different as its lateral lobes are closely spaced and not curved backward (Figs. 8, 9). The other ichnospecies of *Polykampton* show different geometries of the lobes, which are much wider and show a leaf- or petal-like outline (Uchman et al., 2019, 2020). They can easily be distinguished from *P. recurvum* n. isp.

Chondrites recurvus (Brongniart, 1823) shows back-curved branches as in *P. recurvum*, but the branches emanate asymmetrically from a back-curved tunnel or emanate successively one from another. Moreover, their filling does not show any pellets or menisci (Fu, 1991). Generally, this ichnospecies has any median tunnel.

Some similarities are displayed by *Dendrotichnium* Häntzschel, 1975, which is typified by *D. llarenai* (Farrés, 1967) from the Upper Cretaceous of Spain. It has a median tunnel and sparse, lateral, alternating branches, but the branches are never back-curved. All parts of the trace fossil show different proportions than in *Polykampton recurvum* (Fig. 8). Moreover, *Dendrotichnium* is preserved as a convex hypichnion on a turbiditic bed. *D. seilacheri* Kozur, Krainer and Mostler, 1996 from the Permian of Sicily is geometrically somewhat similar. In *D. haentzscheli* (Farrés, 1967), the lateral branches emanate symmetrically from the median tunnel. These representatives of *Dendrotichnium* show no evidence of active filling. Therefore, the branches cannot be treated as lobes, and their relation to *Polykampton* seems to be apparent.

The narrow, almost tubular lobes of *Polykampton recurvum* are somewhat similar to the tubes of *Cladichnus* D'Alessandro and Bromley, 1987. However, *Cladichnus* has no central tunnel, the tubes are branched, and the branching pattern characterized by side offshoots is distinctly different (see Fu, 1991; Wetzel and Uchman, 2013).

Distribution and associated trace fossils

Polykampton recurvum n. isp. occurs in single beds of beige marlstone. In the Campo dei Re section, the bed is 5 cm thick and covered by a 0.7 cm thick layer of gray noncalcareous mudstone (Fig. 2). In the Salata section, *P. recurvum* is present in a 22 cm thick marlstone bed having a 3 cm thick layer of marly siltstone at the base and a 2 cm thick layer of gray marlstone at the top (Fig. 2). After splitting of the beds approximately along a horizontal plane, *P. recurvum* is usually seen on both the lower and upper slit surface. The color contrast

between the burrow filling and the host rock is greater after oxidized.

In the Campo dei Re section, *Polykampton recurvum* occurs in the same bed as *Chondrites intricatus* (Brongniart, 1823) (Figs. 3.1, 4.5, 5.1, 6.2, 6.4, 6.5, 6.7), *Chondrites affinis* (Sternberg, 1833) (Figs. 5.2, 5.5, 6.1, 6.3–6.6), *Palaeophycus* isp. (Figs. 4.5, 5.1, 5.7, 5.8), *Planolites* isp. (Fig. 6.2), *Alcyonidiopsis* isp. (Fig. 5.8), *Phycosiphon geniculatum* (Sternberg, 1833) (Figs. 4.6, 5.7, 6.1–6.5), and possibly *Thalassinoides* isp. (Fig. 6.2). Considering the penetration depth of these trace fossils from the top of the beds, *Polykampton recurvum* occupies a middle tier; only *Chondrites intricatus* more deeply and distinctly crosscuts *P. recurvum*.

In the same bed in the Salata section, *Polykampton recurvum* co-occurs with *Chondrites intricatus* (Fig. 6.7), *Chondrites targionii* (Brongniart, 1828) (Fig. 6.7), *Chondrites affinis*, *Tubulichnium rectum* (Fischer-Ooster, 1858), *Phycosiphon geniculatum*, and a simple, tubular burrow that is 8–9 mm wide (?*Planolites* isp.). The base of the bed contains *Protopalaeodictyon* isp. Here, all the associated burrows descend below *P. recurvum*. In both sections, *P. recurvum* occurs 2–2.5 cm below the top of the marlstone.

The slabs bearing *Polykampton recurvum* from the Fubbiano section contain *Palaeophycus* isp. (Fig. 5.2). The slab with *P. recurvum* from the Rio Ferré section contain *Chondrites intricatus* (Figs. 4.4, 5.3), *Chondrites affinis* (Fig. 4.4), *Phycosiphon geniculatum* (Fig. 5.3), and *Planolites* isp., which was preferentially reworked by the *Chondrites* producer forming “Bandchondriten” sensu Ehrenberg (1942) (Fig. 5.3).

Discussion

The dark mudstone of the lobes and their meniscate structure prove that *Polykampton recurvum* n. isp. is actively filled with mud of similar lithology deposited above the top of the marlstone beds. This implies that the burrow has had an open connection to the seafloor and was formed when the mud had already accumulated as background sediment after deposition of the marly turbidite (Fig. 10). The connection to the seafloor was not detected in the rock although some effort was made for that purpose. The lighter color of the median tunnel suggests that it was an open burrow, which was passively filled in most specimens. However, meniscate filling in one of the specimens (Fig. 5.8) suggests that the tunnel could also be actively filled, probably only in the part that served occasionally as a lobe. It is not clear whether the small pellets in the fill of some specimens were produced by the tracemaker or derived from pelleted sediment on the seafloor. Outward concavity of the menisci in the filling of the lobes shows that the tracemaker entered the sediment laterally from the median tunnel, stuffed the lobes with mud, probably in several repetitive acts,

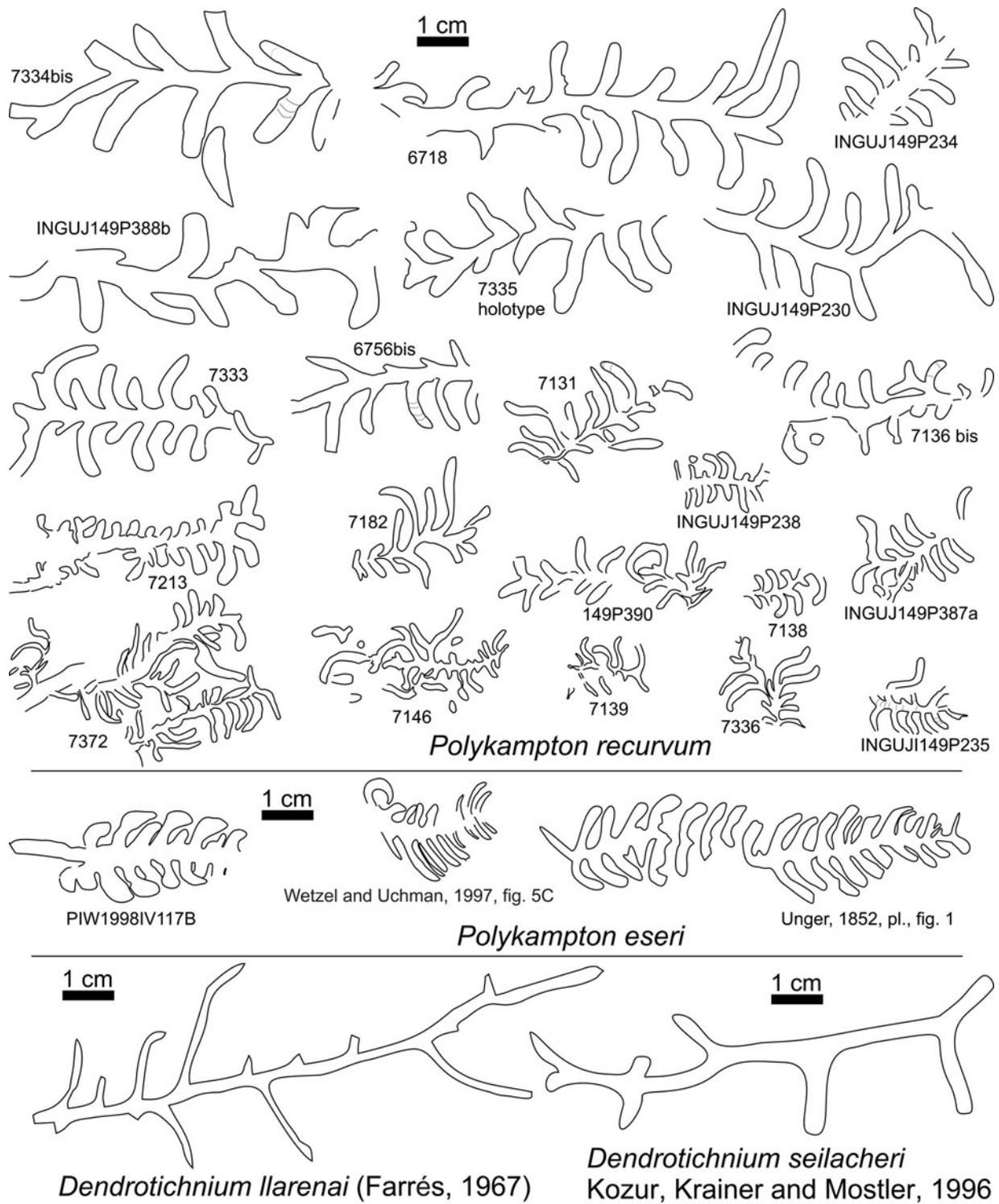


Figure 8. Contours of *Polykampton recurvum* n. isp., *P. eseri*, and *Dendrotichnium* for a comparison of their general shapes.

withdrew to the median tunnel, moved forward, and produced a similar lobe on the other side (Fig. 10). The consistent direction of emanation of the lobes from the median tunnel points to repetitive action initiated from one side of the median tunnel leading to unidirectional propagation of the structure from proximal to distal parts. This suggests a single opening to the

seafloor through which the mud was transported into the tunnel and probes below.

Gentle curvature of the lobes and lobe–tunnel connections suggests a longer, flexible body of a worm-like organism, probably a polychaete, agreeing with prior interpretation of the tracemakers of the other *Polykampton* ichnospecies. Some

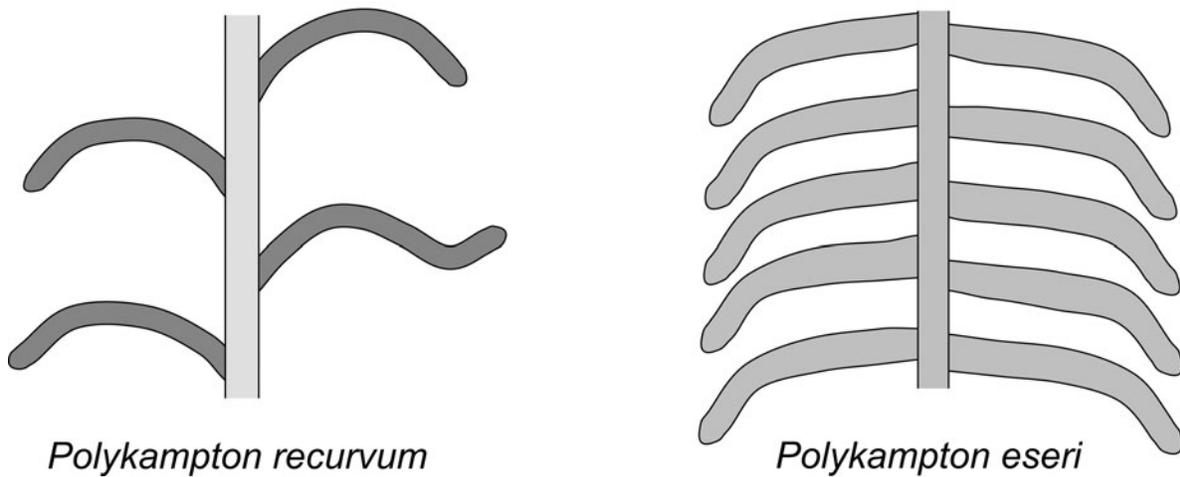


Figure 9. Geometric differences in schematic contours of *Polykampton recurvum* n. isp. and *P. eseri*.

polychaetes are known to drag freshly deposited organic matter into deeper subsurface sediment on the deep-sea floor (Jumars et al., 1990; Levin et al., 1997).

The differences in the width of the lobes within a specimen can be referred to the ontogenetic growth of the tracemaker

during formation of the burrow. The differences in width of the median tunnel and the lobes, and values of other morphometric parameters between specimens (Fig. 7.1, Table 1), point to different sizes of the tracemaker body, which can also be related to the ontogenetic development. This is exemplified by: (1) the distinct

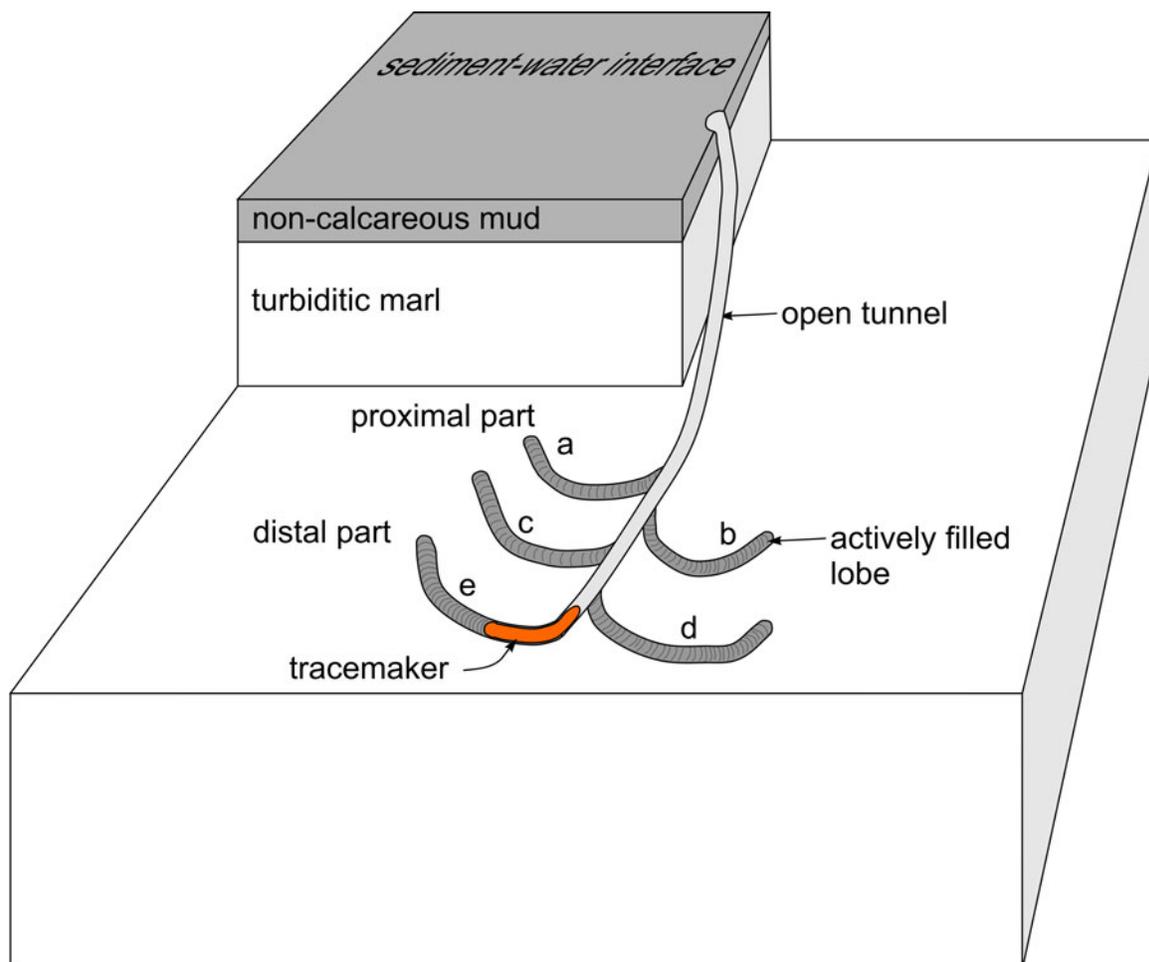


Figure 10. Model of the burrow system and formation of *Polykampton recurvum* n. isp. The letters (a–e) indicate order of production of the lobes.

increase in width of lobes from proximal to distal side of the burrow (Fig. 7.2), (2) the strong ($r = 0.62$) or very strong ($r = 0.71$) positive correlation of minimum/maximum values of lobe width (parameters c_2 , c_1) to those of the median tunnel (a_2/a_1 ; Fig. 7.3, 7.4), (3) the strong positive correlation ($r = 0.68$) between the maximal width of the median tunnel (a_1) and the widest spacing between the lobes (e_1) (Fig. 7.5), (4) the strong positive correlation ($r = 0.67$) between largest width (c_2) and the largest length (d) of the lobes (Fig. 7.6), and (5) the very strong positive correlation ($r = 0.78$) between the maximal (c_1) and minimal (c_2) width of the lobes (Fig. 7.7). The moderate positive correlation ($r = 0.32$) between the smallest and widest spacing between the lobes (e_1/e_2 ; Fig. 7.8) corresponds to the visually poor or moderate regularity of the spacing between the lobes (Figs. 3–6). The width of the narrowest lobe in the proximal part of the burrow may correspond to the width of the body in the initial phase of burrow formation. The maximum width of the tunnel, which usually corresponds to the width of the most distal lobes, reflects the width of the body during the final stage of burrow occupation. This leads to formation of the burrow with a relatively wide median tunnel and lobes whose width increases from proximal to distal parts (Fig. 11). The differences usually do not exceed 50% of the final stage. The ontogenetic development reflected by the differences in size parameters between individual burrows suggests that the tracemaker could form several complete burrows during its life.

According to this interpretation, *Polykampton recurvum* was formed for purposed nutrition. The organic-rich, dark-gray mud was a source of food, especially during periods when little organic matter arrived on the seafloor (= nonbloom times). Therefore, it can be inferred that the lobes served as pantries for food storage. However, there are no traces of reworking of the lobes. This is a repetitive problem in all ichnospecies of *Polykampton*, which can be explained by the possibility that the food can be obtained by ingestion without reworking by means of exoenzymes produced by bacteria, as proposed for the other ichnospecies of *Polykampton* (Uchman and Rattazzi,

2018; Uchman et al., 2019, 2020). Labile organic matter can be released from particulate organic matter by exoenzymes (e.g., Aller and Cochran, 2019) and leached into the main tunnel. As noticed by Boetius (1995) and Chung and King (2001), exoenzymes are concentrated in some polychaete burrows.

Because of the storage of organic-rich mud in *Polykampton recurvum*, it can be treated as another example of the ethological category sequestrichnia, which embraces traces partly or completely filled with sediment utilized as a source of food (Wetzel and Uchman, 2016). Such manner of feeding is typical of the generally oligotrophic deep-sea environments, with a seasonally fluctuating supply of organic matter that is prone to quick oxidization, microbial decomposition, and/or consumption by competitive organisms on the surface (see Druffel et al., 1992). Such predictable supply of organic matter to the deep-sea floor can be seasonal (see Tyler, 1988; Rowe, 2013). It is not excluded that in response to that, the lateral lobes were formed in the same rhythm, as already postulated for *P. cabellae* (Uchman and Rattazzi, 2018). If so, *P. recurvum* was formed at least within 8–24 (mean 13) seasons.

The storage of the organic-matter-rich sediment in a burrow prolongs its nutritional utility. Such behavior is in line with the accelerated diversification of deep-sea trace fossils since the Late Cretaceous and continued through the beginning of the Paleogene because of competition for food (Uchman, 2004). It corresponds to the late phase of the Mesozoic Marine Revolution in the deep sea, establishing the benthic–pelagic coupling pattern, which is already typical of the Modern evolutionary fauna (Buatois et al., 2016).

Location of the beds bearing *Polykampton recurvum* with a depositional system is difficult because of small outcrops and the absence of sedimentary features that would be diagnostic of a common sedimentary model, especially the deep-sea fan with channels, depositional lobes, and associated elements. The abundance of plant detritus in sandy portions of some beds of the sections suggests an extrabasinal origin of the sand, which was supplied from a land by rivers (cf. Zavala et al., 2012). Thick marlstone/mudstone portions in several beds let us suppose that suspension was trapped in some depression. Their material derived probably from a collapsing deeper shelf and slopes. Similar deposition in a deep-sea trench was proposed for the underlying Monte Antola Formation (Fontana et al., 1994). The commonly noncalcareous background mudstones suggest deposition below the calcium compensation depth.

Conclusions

Polykampton recurvum n. isp. is a horizontal, ribbon-like structure composed of a median cylindrical tunnel and sparse, narrow, usually back-curved, lateral lobes, arranged alternately. It occurs 2–3 cm below the top of the Maastrichtian–Paleocene deep-sea turbiditic marlstone beds and is filled with gray mudstone from above. *P. recurvum* was produced by a “worm,” probably a polychaete. The trace fossil belongs to the category sequestrichnia and reflects a generally oligotrophic environment (deep-sea trench below the calcium compensation depth) with periodic supply of organic matter that is quickly oxidized or consumed. Accumulation of organics in the lateral lobes was evidently a source of food for the tracemaker.

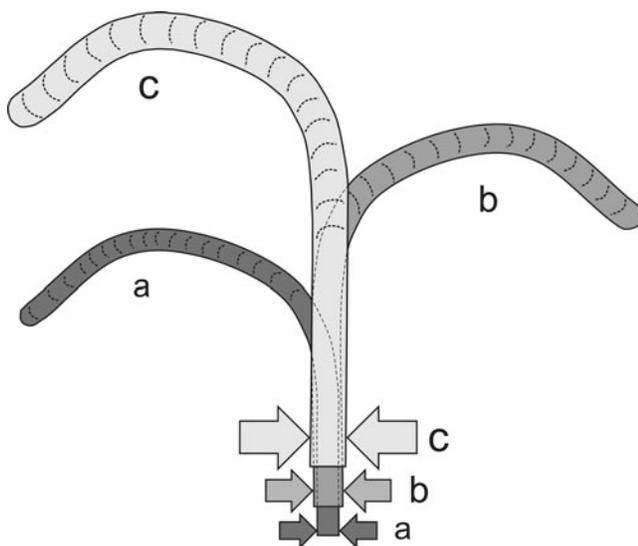


Figure 11. Ontogenetic growth expressed by increasing width of the successive lobes (a, b, c) and width of the median tunnel, which is equal to width of the last lobe.

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References

- Abbate, E., and Sagri, M., 1967, Suddivisioni litostratigrafiche nei calcari ad elmintoidi auctt. della Placca dell'Ebbero-Antola e correlazioni con terreni simili affioranti tra Voghera e Castelnovo ne' Monti (Appennino Settentrionale): *Memorie della Società Geologica Italiana*, v. 6, p. 23–65.
- Aller, R.C., and Cochran, J.K., 2019, The critical role of bioturbation for particle dynamics, priming potential and organic C remineralization in marine sediments: local and basin scales: *Frontiers of Earth Sciences*, v. 7, art. 157, 14 p., <https://doi.org/10.3389/feart.2019.00157>.
- Bellinzona, G., Boni, A., Braga, G., and Marchetti, G., 1971, Note illustrative della Carta Geologica d'Italia in scala 1:100.000, Foglio 71, Voghera: Rome, Servizio Geologico d'Italia, 121 p.
- Boetius, A., 1995, Microbial hydrolytic enzyme activities in deep-sea sediments: Helgoländer Meeresunters., v. 49, p. 177–187.
- Brongniart, A.T., 1823, Observations sur les Fucoïdes: *Société d'Histoire Naturelle de Paris, Mémoire*, v. 1, p. 301–320.
- Brongniart, A.T., 1828, Histoire des Végétaux Fossiles ou Recherches Botaniques et Géologiques sur les Végétaux Renfermés dans les Diverses Couches du Globe, Volume 1: Paris, G. Dufour & E. d'Ocagne, 136 p.
- Buatois, L.A., Carmona, N.B., Curran, H.A., Netto, R.G., Mángano, M.G., and Wetzel, A., 2016, The Mesozoic marine revolution, in Mángano, M.G., and Buatois, L.A., eds., *The Trace-Fossil Record of Major Evolutionary Events, Volume 2: Mesozoic and Cenozoic: Topics in Geobiology*, v. 40, p. 19–134.
- Catanzariti, R., Ellero, A., Levi, N., Ottria, G., and Pandolfi, L., 2007, Calcareous nannofossil biostratigraphy of the Antola Unit succession (Northern Apennines, Italy): new age constraints for the Upper Cretaceous Helminthoid Flysch: *Cretaceous Research*, v. 28, p. 841–860.
- Chung, W.K., and King, G.M., 2001, Isolation, characterization, and polyaromatic hydrocarbon degradation potential of aerobic bacteria from marine macrofaunal burrow sediments and description of *Lutibacterium anuloederans* gen. nov., sp. nov., and *Cycloclasticus spirillensus* sp. nov.: *Applied and Environmental Microbiology*, v. 67, p. 5585–5592.
- D'Alessandro, A., and Bromley, R.G., 1987, Meniscate trace fossils and the *Muensteria-Taenidium* problem: *Palaeontology*, v. 30, p. 743–763.
- Druffel, E.R.M., Williams, P.M., Bauer, J.E., and Ertel, J.R., 1992, Cycling of dissolved and particulate organic matter in the open ocean: *Journal of Geophysical Research*, v. 97, p. 15639–15659.
- Ehrenberg, K., 1942, Über einige Lebensspuren aus dem Oberkreideflysch von Wien und Umgebung: *Paleobiologica*, v. 7 (for 1941), p. 282–313.
- Farrés, F., 1967, Los "Dendroctichium" de España: Notas y Comunicaciones del Instituto Geológico y Minero de España, v. 94, p. 29–36.
- Fischer-Ooster, C., 1858, Die fossilen Fucoïden der Schweizer-Alpen, nebst Erörterungen über deren geologisches Alter: Bern, Huber, 72 p.
- Fontana, D., Spadafora, E., Stefani, C., Stocchi, S., Tateo, F., Villa, G., and Zuffa, G.G., 1994, The Upper Cretaceous Helminthoid Flysch of the northern Apennines: provenance and sedimentation: *Memorie della Società Geologica Italiana*, v. 48, p. 237–250.
- Fu, S., 1991, Funktion, Verhalten, und Einteilung fucoïder und lophoctenider Lebensspuren: *Courier Forschungs-Institut Senckenberg*, v. 135, p. 1–79.
- Gelati, R., and Gnaccolini, M., 1978, I conglomerati della Val Borbera, al margine orientale del bacino Terziario Ligure-Piemontese: *Rivista di Paleontologia e Stratigrafia Italiana*, v. 84, p. 701–728.
- Ghibaud, G., Clari, P., and Perello, M., 1985, Litostratigrafia, sedimentologia ed evoluzione tettonico-sedimentaria dei depositi miocenici del margine sud-orientale del Bacino terziario ligure-piemontese (Valli Borbera, Scrivia e Lemme): *Bollettino della Società Geologica Italiana*, v. 104, p. 349–397.
- Gnaccolini, M., 1988, Osservazioni sui conglomerati Oligocenici affioranti nell'area compresa tra Roccaforte Ligure e Gordona (Alessandria): *Rivista di Paleontologia e Stratigrafia Italiana*, v. 93, p. 521–532.
- Häntzschel, W., 1975, Trace fossils and problematica, in Teichert, C., ed., *Treatise on Invertebrate Paleontology, Part W, Miscellaneous, Supplement I: Boulder and Lawrence, Geological Society of America and University of Kansas Press*, 269 p.
- Jumars, P.A., Mayer, L.M., Deming, J.W., Baross, J.A., and Wheatcroft, R.A., 1990, Deep-sea deposit-feeding strategies suggested by environmental and feeding constraints: *Philosophical Transactions of the Royal Society of London, Series A*, v. 331, p. 85–101.
- Kozur, H.W., Krainer, K., and Mostler, H., 1996, Ichnology and sedimentology of the early Permian deep-water deposits from the Lercara-Roccapalumba area (Western Sicily, Italy): *Facies*, v. 34, p. 123–150.
- Levi, N., Ellero, A., Ottria, G., and Pandolfi, L., 2006, Polyorogenic deformation history recognized at very shallow structural levels: the case of the Antola Unit (Northern Apennine, Italy): *Journal of Structural Geology*, v. 28, p. 1694–1709.
- Levin, L., Blair, N., DeMaster, D., Plaia, G., Fornes, W., Martin, C., and Thomas, C., 1997, Rapid subduction of organic matter by malmanid polychaetes on the North Carolina slope: *Journal of Marine Research*, v. 55, p. 595–611.
- Marroni, M., Mollia, G., Ottria, G., and Pandolfi, L., 2001, Tectono-sedimentary evolution of the External Liguride units (Northern Apennines, Italy): insights in the pre-collisional history of a fossil ocean-continent transition zone: *Geodinamica Acta*, v. 14, p. 307–320.
- Marroni, M., Feroni, A.C., di Biase, D., Ottria, G., Pandolfi, L., and Taini, A., 2002, Polyphase folding at upper structural levels in the Borbera Valley (northern Apennines, Italy): implications for the tectonic evolution of the linkage area between Alps and Apennines: *Compte Rendus Geoscience*, v. 334, p. 565–572.
- Marroni, M., Ottria, G., Pandolfi, G., Catanzariti, R., Bormioli, D., Cucchi, A., and Moletta, G., (in press), Note illustrative della Carta Geologica d'Italia alla scala 1: 50 000, Foglio 196, Cabella Ligure: Roma, Servizio Geologico d'Italia, 178 p.
- Mutti, E., Papani, L., di Biase, D., Davoli, G., Mora, S., Segadelli, S., and Tinterri, R., 1995, Il Bacino Terziario Epimesoalpino e le sue implicazioni sui rapporti tra Alpi ed Appennino: *Memoria di Scienze Geologiche*, v. 47, p. 217–244.
- Ooster, W.-A., 1869, Die organischen Reste der Zoophycos-Schichten der Schweizer Alpen, in Ooster, W.A., and Fischer-Ooster, C.V., eds, *Protozoee Helvetica: Mittheilungen aus dem Berner Museum der Naturgeschichte über merkwürdige Thier- und Pflanzenreste der schweizerischen Vorwelt*, v. 1: Basel, H. Georg, p. 15–35.
- Rowe, G.T., 2013, Seasonality in deep-sea food webs—a tribute to the early works of Paul Tyler: *Deep Sea Research Part II: Topical Studies in Oceanography*, v. 92, p. 9–17.
- Seilacher, A., 2007, *Trace Fossil Analysis*: Berlin, Springer, 226 p.
- Sternberg, G.K., 1833, Versuch einer geognostisch-botanischen Darstellung der Flora der Vorwelt. IV Heft: Regensburg, C.E. Brenck, 48 p.
- Tyler, P.A., 1988, Seasonality in the deep sea: *Oceanography and Marine Biology: An Annual Review*, v. 26, p. 227–258.
- Uchman, A., 1999, Ichnology of the Rhenodanubian Flysch (Lower Cretaceous–Eocene) in Austria and Germany: *Beringeria*, v. 25, p. 65–171.
- Uchman, A., 2004, Phanerozoic history of deep-sea trace fossils, in McIlroy, D., ed., *The application of ichnology to palaeoenvironmental and stratigraphic analysis*: Geological Society of London Special Publication, v. 228, p. 125–139.
- Uchman, A., 2007, Trace fossils of the Pagliaro Formation (Paleocene) in the North Apennines, Italy: *Beringeria*, v. 37, p. 217–237.
- Uchman, A., and Rattazzi, B., 2018, The trace fossil *Polykampton cabellae* isp. nov. from the Pagliaro Formation (Paleocene), Northern Apennines, Italy: a record of nutritional sediment sequestration by a deep sea invertebrate: *Ichnos*, v. 25, p. 1–10.
- Uchman, A., Wetzel, A., and Rattazzi, B., 2019, Alternating stripmining and sequestration in deep-sea sediments: the trace fossil *Polykampton*—an ecological and ichnotaxonomic evaluation: *Palaeontologia Electronica*, 22.2.21A, 18 p., <https://doi.org/10.26879/930>
- Uchman, A., Lebanidze, Z., Beridze, T., Kobakhidze, N., Lobzhanidze, K., Khutsishvili, S., Chagelishvili, R., Makadze, D., Koiava, K., and Khundadze, N., 2020, Abundant trace fossil *Polykampton* in Palaeogene deep-sea flysch deposits of the Lesser Caucasus in Georgia: palaeoecological and palaeoenvironmental implications: *Palaeogeography, Palaeoclimatology, Palaeoecology*, v. 558, n. 109958, <https://doi.org/10.1016/j.palaeo.2020.109958>
- Unger, F., 1850, Genera et species plantarum fossilium: Vindobonae, Wilhelm Braumüller, 627 p.
- Unger, F., 1852, *Iconographia plantarum fossilium. Abbildungen und Beschreibung fossiler Pflanzen*: Denkschriften der kaiserlichen akademie der Wissenschaften. Mathematisch-naturwissenschaftlichen Classe, v. 4, p. 73–118.
- Wetzel, A., and Uchman, A., 1997, Ichnology of deep sea fan overbank deposits of the Ganei Slat (Eocene, Switzerland)—a classical flysch trace fossil locality studied first by Oswald Heer: *Ichnos*, v. 5, p. 139–162.
- Wetzel, A., and Uchman, A., 2013, *Cladichnus parallelum* isp. nov.—a new mid- to deep-tier feeding burrow system: *Ichnos*, v. 20, p. 120–128.

Wetzel, A., and Uchman, A., 2016, Sequestrichnia—a new ethological category of trace fossils in oligotrophic deep-sea environments, in Baucon, A., Neto de Carvalho, and Rodrigues, J., eds., *Ichnia 2016, Abstract Book: Castelo Branco, UNESCO Geopark Naturejo and International Ichnological Association*, p. 190.

Zavala, C., Arcuri, M., and Blanco Valiente, L., 2012, The importance of plant remains as diagnostic criteria for the recognition of ancient hyperpycnites: *Revue de Paléobiologie*, v. 11, p. 457–469.

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