

# Bryozoan fauna from the Reynales Formation (lower Silurian, Aeronian) of New York, USA

Andrej Ernst,<sup>1</sup> Carlton E. Brett,<sup>2</sup> and Mark A. Wilson<sup>3</sup>

<sup>1</sup>Institut für Geologie, Universität Hamburg, Bundesstr. 55, D-20146 Hamburg, Germany <[Andrej.Ernst@uni-hamburg.de](mailto:Andrej.Ernst@uni-hamburg.de)>

<sup>2</sup>Department of Geology, University of Cincinnati, Cincinnati, Ohio 45221-0013, USA <[brettce@ucmail.uc.edu](mailto:brettce@ucmail.uc.edu)>

<sup>3</sup>Department of Earth Sciences, The College of Wooster, Wooster, Ohio 44691, USA <[mwilson@wooster.edu](mailto:mwilson@wooster.edu)>

**Abstract.**—Thirteen bryozoan species are described from the Brewer Dock (Hickory Corners) Member of the Reynales Formation (lower Silurian, Aeronian) at the locality Hickory Corners in western New York, USA. Three species are new: trepostomes *Homotrypa niagarensis* n. sp. and *Leioclema adsuetum* n. sp. and the rhabdomesine cryptostome *Moyerella parva* n. sp. Only one species, *Hennigopora apta* Perry and Hattin, 1960, developed obligatory encrusting colonies whereas the others produced erect ramose colonies of various thicknesses and shapes: cylindrical, branched, and lenticular. Bryozoans display high abundance and richness within the rock. This fauna is characteristic of a moderately agitated environment with a stable substrate. The identified species reveal paleobiogeographic connections to other Silurian localities of New York as well as Ohio and Indiana (USA) and Anticosti (Canada).

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## Introduction

Silurian rocks exposed in classic localities of northeastern United States contain abundant fossils, including bryozoans. However, the last major studies of Silurian Bryozoa from these localities were conducted more than 100 years ago. Whereas bryozoans from the Silurian of Ohio and Indiana were described in an impressive series of monographs and journal articles (e.g., Hall and Whitfield, 1875; Hall, 1876, 1882; Foerste, 1887, 1888, 1893, 1919; Perry and Hattin, 1960), bryozoan faunas of western New York received little attention (e.g., Hall, 1852; Bassler, 1906). In addition, the majority of literature on American Silurian bryozoans predates the present-day standards of taxonomic descriptions.

Silurian bryozoans are known from many local areas worldwide; however, their paleontological record is not as remarkable as for Ordovician and Devonian faunas. Tuckey (1990) stated that a low level of endemism exists for Silurian bryozoans and postulated a correlation between provinciality and bryozoan diversity. Controversially, McCoy and Anstey (2010) concluded that bryozoans showed a high level of endemism in the Silurian, much higher than other organism groups. Buttler et al. (2013) showed that the provinciality was heterogeneous during the Silurian. During the Llandovery, bryozoans began to show distinct provincialism; this declined during the Wenlock, only to reemerge during the Ludlow.

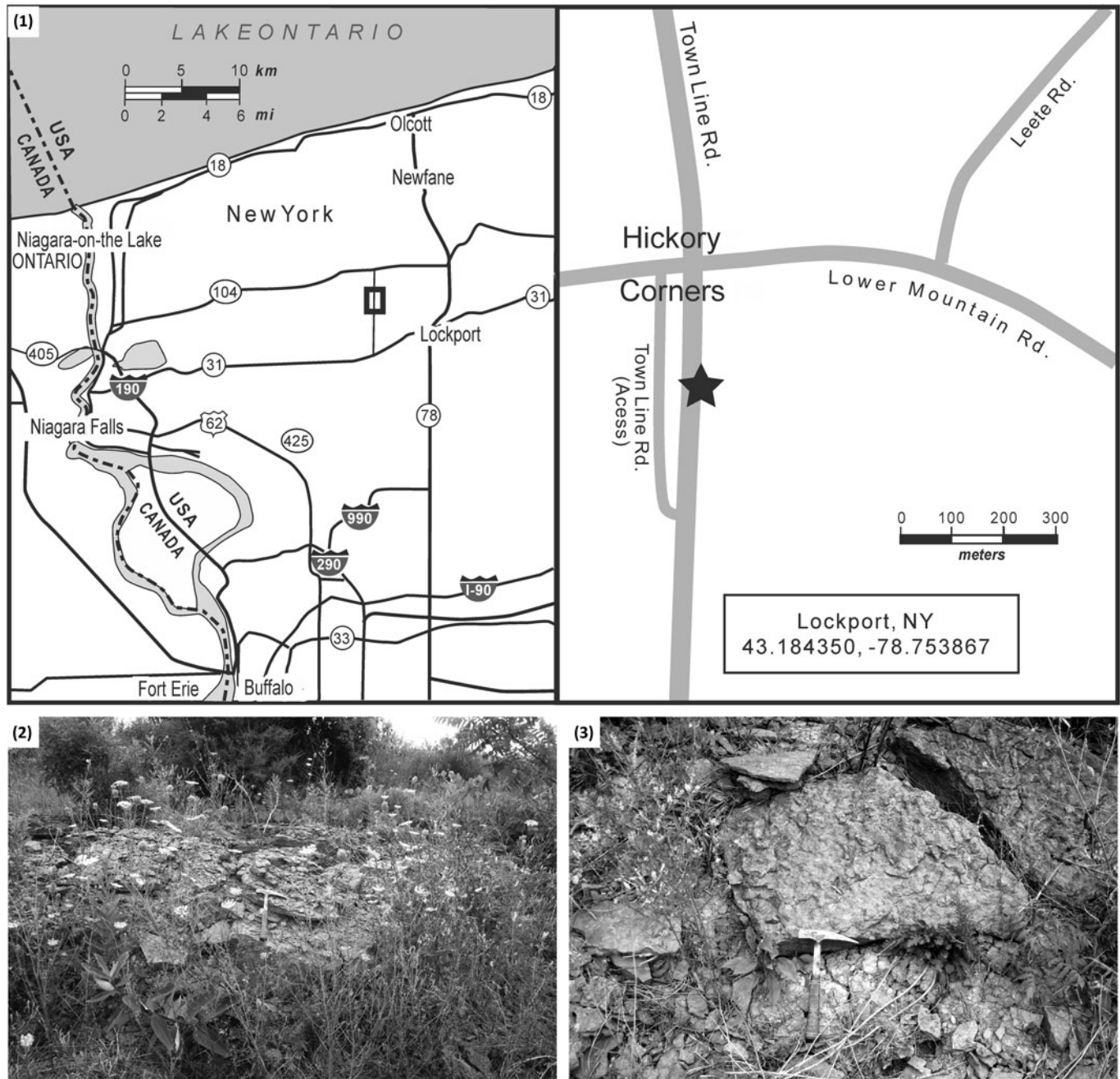
Bryozoans are a very important fossil group. Because their skeletons consist of carbonate material (usually low-magnesium calcite in Paleozoic clades), bryozoans show high potential for preservation. Because of their modular organization, bryozoan colonies can be identified even from small fragments. The

wide distribution of bryozoans in marine habitats and their ecological flexibility make them important for use in various fields of the life sciences. The present paper documents a little-known Aeronian bryozoan fauna from New York. In all, 13 species, including three new species, are described, and the paleoecologic and biogeographic implications of this fauna are discussed.

## Geological and paleontological setting

This bryozoan fauna derives from the Reynales Formation (lower Silurian, Llandovery, Aeronian) at Hickory Corners, Lockport Junction Road (Route 93), near the city of Lockport, Niagara County, New York (Fig. 1). The studied section comprises about two meters of interbedded shale and limestones represented by biosparite (grainstones) and biomicrite (wackestones and packstones) with a carbonate hardground capping the unit (Fig. 2). Kilgour (1963) named these rocks the Hickory Corners Member of the Reynales Formation. LoDuca and Brett (1994) demonstrated that the unit at this outcrop is actually a composite western extension of the Brewer Dock and Wallington members of the Reynales of the Rochester area (Gillette, 1947) and that those names would have priority over Hickory Corners Member. Nonetheless, the boundary between these units is difficult to recognize at Lockport. The name Hickory Corners is officially recognized by U.S. Geological Survey GEOLEX, and this name will be used herein.

The Reynales Formation has long been known to be of mid-Llandovery and probably Aeronian age (Berry and Boucot, 1970). Recently, Waid and Over (2015) discovered conodonts in the lower Reynales Formation (basal Budd Road phosphate bed of LoDuca and Brett, 1994) from Budd Road about 2.5

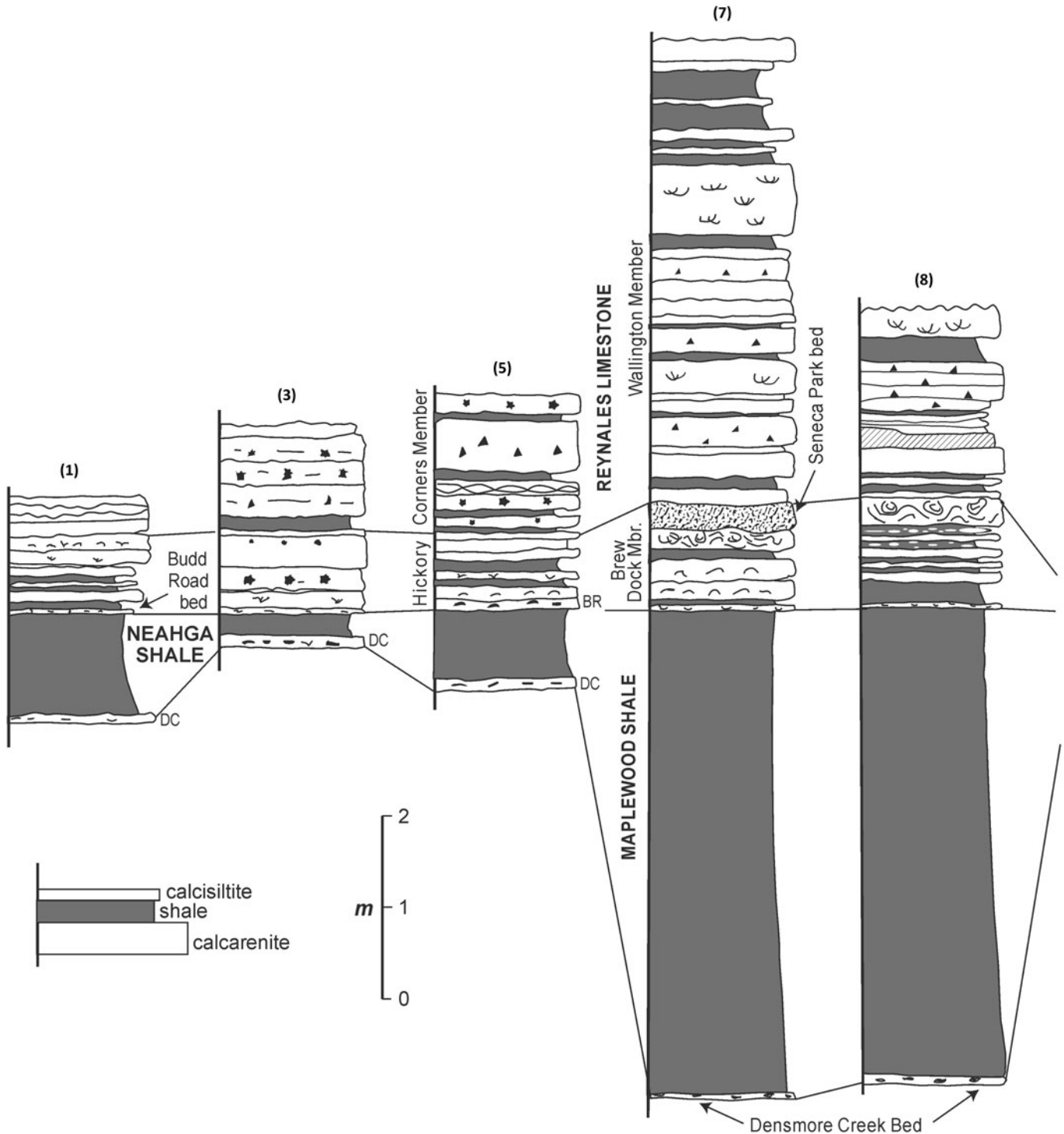


**Figure 1.** (1) Locality map showing position of the Hickory Corners (Lockport Junction Road, Route 93) outcrop; (2, 3) external view of the outcrop showing limestones of the Reynales Formation.

km west of Hickory Corners, representing the *Pranognathus tenuis* Zone (lower to middle Aeronian). The Reynales Formation is at the top of Silurian third-order sequence S-II of the lower Clinton Group (Brett et al., 1990; Cramer et al., 2011). In western New York, the Hickory Corners Member interval is represented by an erosional remnant, which pinches out westward in Ontario beneath the Telychian Merrittton Formation (Kilgour, 1963; Fig. 3). It passes eastwardly into the Brewer Dock and lower portion of the Wallington Member of the Reynales Formation, and that in turn a distinctive fossiliferous hematitic ironstone, the Furnaceville Member, is associated with the basal Brewer Dock Member at the Genesee Gorge in Rochester, New York

(Fig. 3). No hematitic bed occurs in the equivalent lower Hickory Corners Member in Niagara County (Fig. 3).

The Reynales Formation was formed in the northern portion of the elongated Appalachian Foreland Basin during the early Silurian Aeronian Age. Deposits extended northeast-southwest in the New York region. During Reynales deposition, the basin center is inferred to have lain to the west of the study area, but western facies have been largely removed by regional erosion in the later Telychian. To the east, the Reynales Formation shows a gradual change to somewhat shallower facies near Rochester, New York (LoDuca and Brett, 1994; Eckert and Brett, 2001). This portion of the Laurentian continent (ancestral

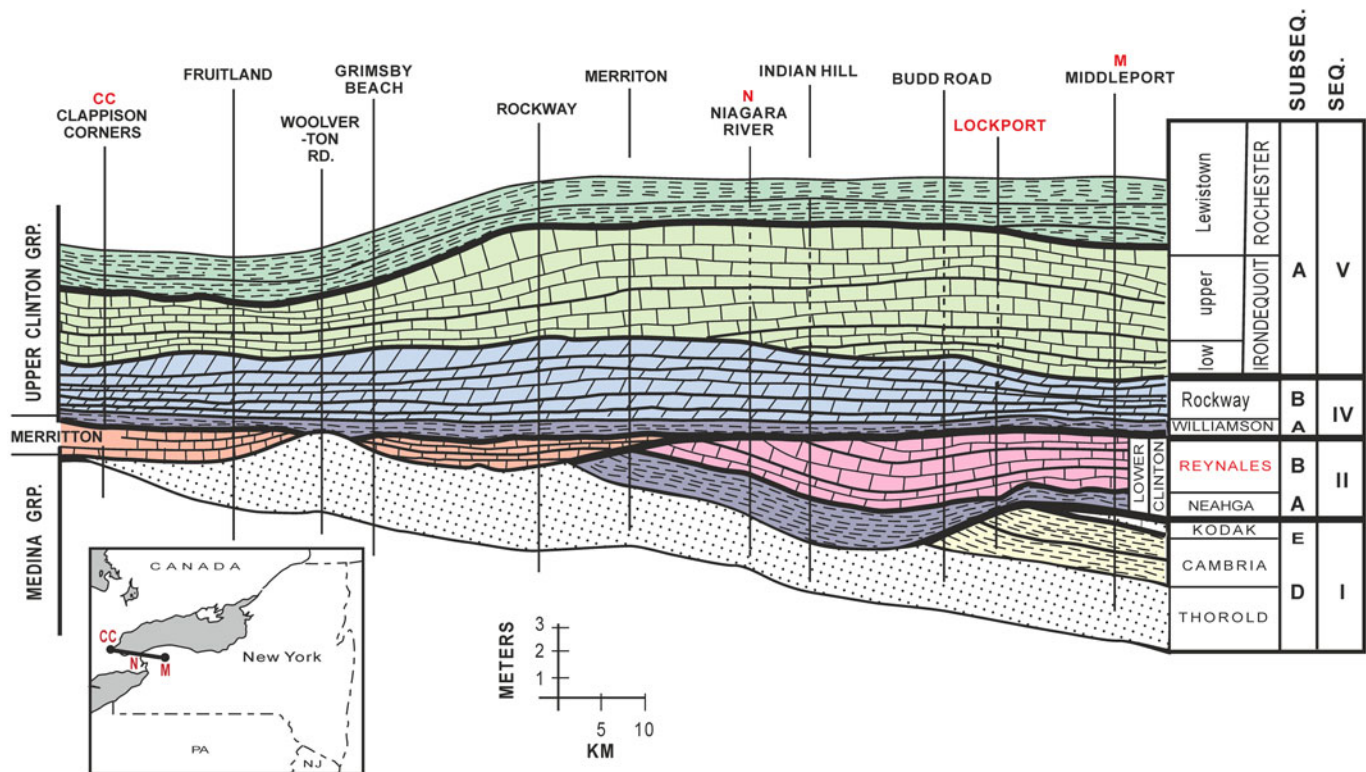


**Figure 2.** Stratigraphy of the Reynales Formation from west (left) to east (right) along the Niagara Escarpment, about 145 km in western New York; locations of the sections are as follows: 1 = Niagara Gorge at Lewiston, Niagara County, New York (43.1479°N, 79.0410°W); 3 = Budd Road, Cambria, Niagara County, New York (43.1843°N, 78.7804°W); 5 = study section: Hickory Corners, Lockport Junction Road (NY 93); Pekin/Lockport, Niagara County, New York (43.1844°N, 78.7539°W); 7 = Genesee River Gorge at Seth Green Drive, Rochester, Monroe County, New York (43.1861°N, 77.6247°W); 8 = Glen Edyth Drive, Webster, Monroe County, New York (43.2044°N, 77.5137°W); these and the omitted localities are discussed in LoDuca and Brett (1994), from which this figure is adapted.

North America) was subtropical, situated about 30° south of the equator (Witzke, 1990; Scotese, 2001; Blakey, 2005).

The Hickory Corners Member comprises a thin interval of interbedded, sparsely fossiliferous greenish gray shale and fossiliferous wackestones and packstones with minor crinoidal grainstones near the top. Beds are thin and commonly lenticular;

some show evidence of grading suggestive of storm processing of skeletal material and silty carbonate muds. The basal few centimeters (Budd Road beds) contain abundant shiny black phosphatic nodules suggestive of very slow rates of sediment and prolonged reworking and time averaging, as well as a source of phosphatic material; this is possibly associated with the



**Figure 3.** Regional cross section of the condensed interval of the Silurian rocks in western New York; note the several sequence-bounding unconformities and westward taper and pinchout of the Aeronian Reynales Formation. Sequence II, subsequence B, the study interval, is highlighted in pink. CC, N, and M refer to positions marked on the cross section on the inset map. Modified from LoDuca and Brett (1994).

deposition of the Furnaceville hematite bed, which contains abundant bryozoan fragments (*Phaenopora?* and small ramose bryozoans) coated with hematite. A majority of the brachiopods are disarticulated whole valves without much fragmentation, and the bryozoans are partially intact, branched colonies. This evidence suggests a predominantly low-energy setting with frequent storm agitation and an overall low rate of sediment accumulation. However, layers about 30–50 cm above the base of the Hickory Corners yield evidence of rapid redeposition; these include rare obrution beds that enclose complete crinoids buried in thin mud layers (Eckert and Brett, 2001). The diverse fauna and primarily matrix-supported limestones indicate that the Reynales Formation in this locality represents the shallow (30–60 m depth) outer shelf environment of Benthic Assemblage 4 of Brett et al. (1993).

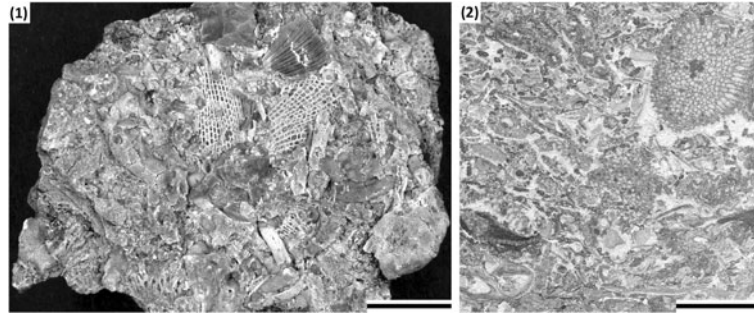
*Associated fauna.*—The Hickory Corners Member is rich in bryozoans, crinoids, bivalves, gastropods, brachiopods, and trilobites (Fig. 4). Small, solitary rugose corals (*Enterolasma?* sp.) are common in this limestone. Brachiopods are relatively diverse and include *Hyattidina*, *Platystrophia*, *Dalejina*, *Coolinia*, and rare *Eocoelia*; disarticulated sclera of the trilobites *Liocalymene* and *Encrinurus* are relatively common in the Hickory Corners Member.

*Eocoelia*, often considered to be a nearshore indicator, is typical of the ‘*Eocoelia* Community’ (Ziegler, 1965) or benthic assemblage (BA) 2 of Boucot (1975; see Boucot and Lawson, 1999 for numerous examples). However, this brachiopod is rare in the Niagara County area. The upper portion of the Hickory

Corners Member passes eastward (up-ramp) into the Wallington Limestone, which is rich in robust pentamerid brachiopods (*Pentamerus oblongus* Sowerby, 1839), as well as favositids and small stromatoporoids, a typical BA3 assemblage (Boucot, 1975; Boucot and Lawson, 1999). Overall, the moderate diversity of brachiopod, bryozoan, and echinoderm fauna of the Hickory Corners Member is indicative of a BA3 to BA4 position.

The Reynales Formation in Niagara County, including the Hickory Corners study locality, has yielded a small but distinctive crinoid fauna (Eckert and Brett, 2001), including the small disparid crinoid *Haptocrinus calvatus* Eckert and Brett, 2001, long known for its distinctive pentalobate columnals, which make up a high proportion of the Hickory Corners grainstones. Hall (1852) figured these under an incompletely known species ‘*Glyptocrinus plumosus*.’ These small crinoids are associated with the coiled columns of perhaps three species of *Eomyelodactylus*, notably *E. sparteus* Eckert, 1990, the camerate *Dynamocrinus robustus* Eckert and Brett, 2001, and the flexible crinoid *Prolixocrinus anellus* Eckert and Brett, 2001.

The Hickory Corners faunas, especially the crinoids, show affinities to those described from the uppermost Brassfield Formation (‘Red Brassfield’) in Ohio and Kentucky (e.g., Ausich, 1984, 1986a, b; Ausich and Dravage, 1988). On the basis of conodonts and the brachiopods *Microcardinalia* and *Triplecia*, this unit has been suggested to be of mid-Aeronian age and thus equivalent to the Oldham Limestone of Kentucky and ‘Packer Shell’ of the Ohio Subsurface, both of which are correlated with the Reynales Formation in New York. All of these units yield moderately abundant ramose and reticulate bryozoans



**Figure 4.** (1) Surface of fossil-rich limestone of the Reynales Formation; (2) thin section across bedding. (1) Scale bar = 10 mm; (2) scale bar = 5 mm.

that have not received recent study (see Foerste, 1887, 1893, 1919) and warrant further comparative work.

### Materials and methods

The bryozoans described here were collected from a small weathered roadcut outcrop approximately 2 m high on the east side of State Route 93 (Lockport Junction Road) just south of the Lower Mountain Road overpass in the hamlet of Hickory Corners, part of the town of Cambria, Niagara County, New York, USA (43.18464°N, 78.75394°W). Here the Reynales Formation (lower Silurian, Aeronian) is exposed for several meters along the highway. Most material was obtained from loose slabs that can only be from the Hickory Corners Member; these are mainly from above the basal Budd Road phosphatic beds, but the exact horizon could not be determined.

Bryozoans were studied in thin section using a binocular microscope in transmitted light. In total, 87 thin sections were prepared from rock samples. Morphological character terminology is partly adopted from Anstey and Perry (1970) for trepostomes and cystoporates, Hageman (1991) for fenestrates, and Hageman (1993) for cryptostomes. The spacing of structures is measured as the distance between their centers. Statistics were summarized using arithmetic mean, sample standard deviation, coefficient of variation, and minimum and maximum values.

*Repositories and institutional abbreviations.*—The studied material is deposited at the Senckenberg Museum, Frankfurt am Main, Germany (SMF). Other repositories include the National Museum of Natural History (formerly United States National Museum), Paleobiology Department (USNM PAL), Washington, D.C., USA, and the Geological Survey of Canada (GSC), Ottawa, Canada.

### Systematic paleontology

Phylum Bryozoa Ehrenberg, 1831  
 Class Stenolaemata Borg, 1926  
 Order Cyclostomata Busk, 1852  
 Family Diploclemaeidae Gorjunova, 1992  
 Genus *Diploclema* Ulrich in Miller, 1889

1889 *Diploclema* Ulrich in Miller, p. 300.  
 1890 *Diploclema*; Ulrich, p. 368.  
 1906 *Diploclema*; Bassler, p. 17.

1964 *Diploclema*; Ross, p. 28.  
 1971 *Diploclema*; Kopajevich, p. 253.  
 1975 *Diploclema*; Brood, p. 109.  
 2012 *Diploclema*; Ernst and Carrera, p. 722.

*Type species.*—*Diploclema trentonense* Ulrich in Miller, 1889. Middle Ordovician ('Trenton Group'), USA

*Diagnosis (emended).*—Branched dichotomous colonies. Autozoecia pyriform, growing in biradial pattern from the median lamina through simple exterior skeletal wall, separated by interzoecial space in endozone. In some species, interzoecial space protrudes into the exozone between autozoecia giving an impression of tubes. Autozoecial apertures rounded to oval, with low peristomes, opening distally on all sides of subcylindrical stems, arranged in quincunx order on the colony surface, surrounded by peripheral structures and nodes. The exterior skeletal walls thin, continuous except for autozoecial apertures, covering both autozoecial chambers and small portions of colony-wide extrazoecial space, consisting of prismatic material. Autozoecial diaphragms rarely occurring. Interior autozoecial walls thickly laminated, with distinct zooecial boundaries, often undulating in proximal parts of autozoecia. Communication pores absent. Pseudopores absent. Heterozooecia not known.

*Occurrence.*—Middle Ordovician of United States. Upper Ordovician of western Argentina. Lower to middle Silurian of United States, Canada, and Europe.

*Remarks.*—The genus *Diploclema* Ulrich in Miller, 1889 is unique among cyclostome bryozoans because of its exterior wall, which has a prismatic structure and is completely different from the laminated interior wall. *Diploclema* can be compared with *Kukersella* Bassler, 1911 in possessing similar colony form and autozoecial shape, but *Diploclema* lacks axial tubes, communication pores, and pseudopores. Furthermore, the microstructure of the frontal walls of *Kukersella* is finely laminated whereas the frontal wall of *Diploclema* is prismatic.

*Diploclema argutum* Bassler, 1906  
 Figures 5.1–5.9, 6.1–6.3; Table 1

1906 *Diploclema sparsum* var. *argutum* Bassler, p. 17, pl. 5, figs. 8, 9.

1971 *Diploclema regulariforme* Kopajevich, 1971, p. 121, fig. 3.

*Holotype*.—USNM PAL 35557. Rochester Shale, Wenlock, lower Silurian; New York, USA.

*Occurrence*.—Lower Silurian, Llandovery-Wenlock of USA (New York). Lower Silurian, Wenlock of Estonia.

*Description*.—Branched dichotomous colonies, cylindrical and belt shaped, bifoliate, arising from encrusting bases. Cylindrical branches 0.30–0.46 mm in diameter, belt shaped 0.37–0.43 mm thick and 0.75–1.43 mm wide. Autozoecia drop shaped, narrow proximally and broadened distally, 0.40–0.60 mm long and 0.13–0.20 mm wide, separated by interzoecial space in endozone. Interzoecial space protruding to the exterior wall between autozoecia producing rounded separations 0.04–0.08 mm in diameter. Autozoecial apertures oval, 0.07–0.10 mm wide, spaced 0.34–0.44 mm from center to center longitudinally. Interior autozoecial walls thickly laminated, undulating near the surface, with distinct zoecial boundaries, 0.013–0.040 mm thick. Exterior zoecial walls granular, 0.005–0.008 mm thick.

*Materials*.—SMF 23.318–SMF 23.376.

*Remarks*.—Bassler (1906, p. 18, pl. 5, figs 8, 9) described “a distinct pore in front of zoecial aperture.” These pores were sometimes regarded as heterozoecia. Kopajevich (1971, p. 120, text-figs. 1, 2) described ‘upper’ and ‘lower’ partitions in the autozoecia of *Diploclema*. This model implies zoecia with long peristomes and arched zoecia. A shallow section that cuts the peristome (upper partition) and the arched part of the zoecium (lower partition) produces the appearance of heterozoecia. A similar model was suggested by Brood (1975, p. 64, fig. 12). He suggested the existence of a “vertically placed tabula, which divides the zoecium in two compartments.” The outer compartment is supposed to have contact with the surface, whereas the inner compartment would be connected by pores to the outer one. However, such a morphology is not obvious from observation of the presently described abundant material from the Silurian of North America. The morphology of *Diploclema* is difficult to describe because its outer wall is extremely thin and cannot be observed in tangential sections. Transverse and longitudinal sections show that zoecia are rather flat, and they have neither distinct peristomes nor compartments (Fig. 6.1). Any openings of the ‘heterozoecia’ or ‘pores’ could not be observed in the tangential section. *Diploclema* has a kind of an interzoecial space, which separates autozoecia in endozone, and in distal parts it protrudes from the exozone and reaches the exterior wall (but does not protrude into it). This space is filled by calcite spar, which means that it had no direct contact with seawater. By contrast, autozoecial space is filled by micrite, assuming contact with the sediment via the aperture (after death and decay of the bryozoan). The impression of ‘partitions’ or ‘compartments’ inside of the autozoecia of *Diploclema* appears in oblique sections (Fig. 5.4), whereas strictly longitudinal sections cut only the autozoecia (Fig. 6.1).

*Diploclema argutum* Bassler, 1906 is identical to *D. regulariforme* Kopajevich, 1971 from the Silurian (Wenlock) of Estonia. The two species coincide in their size (e.g., aperture width 0.07–0.10 mm versus 0.05–0.11 mm in *D. regulariforme*; autozoecial width 0.13–0.20 mm versus 0.13–0.18 mm in *D. regulariforme*; autozoecial length 0.40–0.60 mm versus 0.49–0.63 mm in *D. regulariforme*). *Diploclema argutum* is similar to *D. sparsum* (Hall, 1852) from the Silurian (Wenlock) of New York but differs from the latter in coarser branches that are often flat and contain mesotheca instead of median axis.

Superorder Palaeostomata Ma, Buttler, and Taylor, 2014  
 Order Cystoporata Astrova, 1964  
 Family Fistuliporidae Ulrich, 1882  
 Genus *Cheilotrypa* Ulrich, 1884  
 [= *Chilotrypa* Miller, 1889, incorrect subsequent spelling]

*Type species*.—*Cheilotrypa hispida* Ulrich, 1884. Carboniferous, Mississippian, Viséan-Serpukhovian; USA.

*Cheilotrypa* aff. *C. variolata* (Hall, 1876)  
 Figures 6.4–6.8, 7.1–7.5; Table 2

1876 *Trematopora variolata* Hall, p. 113, pl. 11, figs. 9, 10.

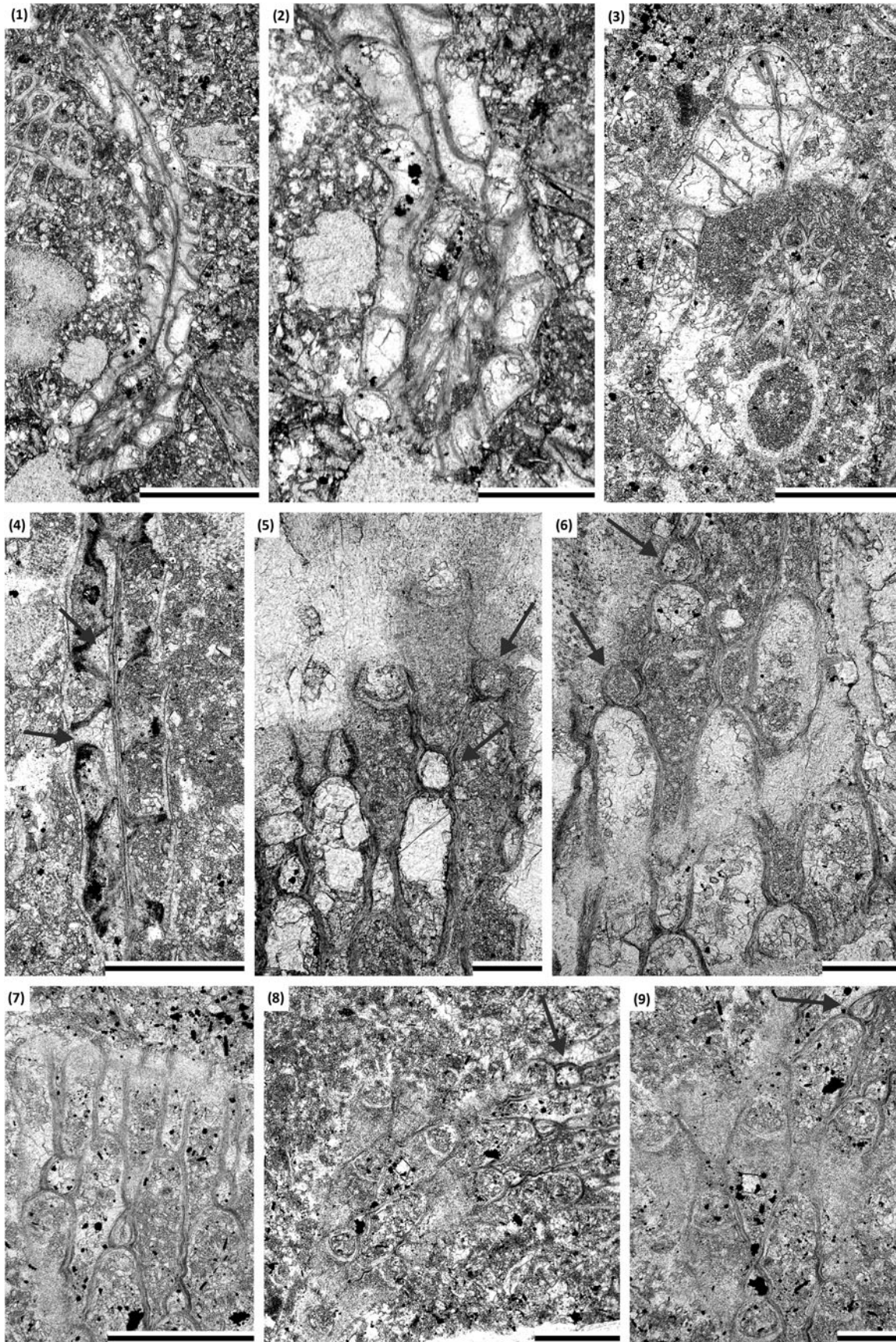
1882 *Trematopora variolata*; Hall, p. 234, pl. 10, figs. 9, 10.

*Occurrence*.—Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA.

*Description*.—Cylindrical colonies formed by encrusting unpreserved objects, 1.25–2.50 mm in diameter, with a lumen 0.42–1.35 mm in diameter. Encrusting sheets 0.26–1.10 mm in thickness, locally budding ‘base to base’ (Fig. 6.7, 6.8). Encrusted objects represent apparently dichotomizing rigid cylindrical objects (algae?) with regular narrow portions (Fig. 6.6). Autozoecia growing from 0.005–0.010 mm thick epitheca, bending sharply at their bases toward colony surface. Autozoecial apertures circular to oval. Basal diaphragms rare to absent. Lunaria well developed, horseshoe shaped. Vesicles abundant, five to eight surrounding each autozoecial aperture, irregularly shaped, medium in size, often large at the base, polygonal in tangential section, box-like to hemispheric, with plane or concave roofs in exozone. Autozoecial walls granular, 0.003–0.005 mm thick. Extensive extrazoecial material in exozone developed, 0.20–0.33 mm thick. Maculae consist of vesicular skeleton covered by laminated calcitic material, 0.38–0.70 mm in diameter.

*Materials*.—SMF 23.377–SMF 23.461.

*Remarks*.—Three species of *Cheilotrypa* are known from the Silurian of North America: *C. varia* (Hall, 1876), *C. variolata* (Hall, 1876), and *C. ostiolata* (Hall, 1852) (all from the Wenlock of Indiana and New York). Whereas the former two species were described without use of thin sections, *Cheilotrypa ostiolata* was identified and described internally by Perry and



Hattin (1960) from the Osgood Formation (Wenlock) of Indiana. According to the descriptions and illustrations of *C. varia* and *C. variolata* in Hall (1882), the present species appears most similar to *C. variolata*. Hall (1882, p. 234) gives branch diameters of 2–7 mm and aperture diameters of 0.25 mm for *C. varia* and branch diameters of 1–1.5 mm and aperture width of 0.20 mm for *C. variolata*. The aperture width is still larger than in the present species (0.09–0.15 mm). *Cheilotrypa* sp. differs from *C. ostiolata* from the Silurian of New York in smaller autozoecia (autozoecia width 0.09–0.15 mm versus 0.15–0.20 mm in *Cheilotrypa ostiolata*).

Family Xenotrypidae Utgaard, 1983  
Genus *Hennigopora* Bassler, 1952

*Type species*.—*Callopora florida* Hall, 1852. Niagaran Group, Rochester Shale, Sheinwoodian, Wenlock, Silurian; Lockport, New York, USA.

*Hennigopora apta* Perry and Hattin, 1960  
Figure 7.6–7.9; Table 3

1960 *Hennigopora apta* Perry and Hattin, p. 704, pl. 88, figs. 1, 3, 4.

*Holotype*.—Specimen 5572 (Indiana University Paleontological Collection). Osgood Formation (Telychian-Sheinwoodian), Silurian; Indiana, USA.

*Occurrence*.—Osgood Formation (Telychian-Sheinwoodian), Silurian; Indiana, USA. Reynales Formation, Hickory Corners Member, Aeronian, Llandoverly, lower Silurian; Hickory Corners, New York, USA.

*Description*.—Encrusting colony, 1.25 mm thick. Autozoecia growing from thin epitheca, bending in the early exozone to the colony surface. Autozoecial diaphragms rare; straight, thin. Autozoecial apertures rounded polygonal, strongly petaloid due to indenting acanthostyles. Vesicles generally large, separating autozoecia in one row, 8–11 surrounding each autozoecial aperture, with rounded roofs, polygonal in tangential section. Acanthostyles abundant, relatively large, often deeply indenting autozoecial chambers, three to five surrounding each autozoecial aperture. Autozoecial walls displaying obscure granular microstructure, 0.005–0.015 mm thick. Maculae not observed.

*Materials*.—SMF 23.462–SMF 23.469.

*Remarks*.—*Hennigopora apta* Perry and Hattin, 1960 differs from *H. florida* (Hall, 1852) in having smaller autozoecial apertures (0.17–0.25 mm versus 0.28–0.35 mm in *H. florida*).

Order Trepostomata Ulrich, 1882  
Suborder Halloporina Astrova, 1965  
Family Monticuliporidae Nicholson, 1881  
Genus *Homotrypa* Ulrich, 1882  
[= *Homotrypella* Ulrich, 1882]

*Type species*.—*Homotrypa curvata* Ulrich, 1882. Upper Ordovician (Cincinnatian); North America.

*Homotrypa niagarensis* new species  
Figures 8.1–8.9, 9.1; Table 4

*Holotype*.—SMF 23.470–SMF 23.473 (four thin sections), SMF 23.474 (rock sample).

*Paratypes*.—SMF 23.475–SMF 23.502.

*Diagnosis*.—Ramosely branched colonies with narrow exozones; autozoecia long, bending at intermediate angles in exozone, having rounded polygonal apertures; basal diaphragms in exozone; cystiphagms large, abundant in exozone; mesozooecia rare to common, small; acanthostyles rare to common, large; autozoecial walls thick, laminated with distinct medial lining.

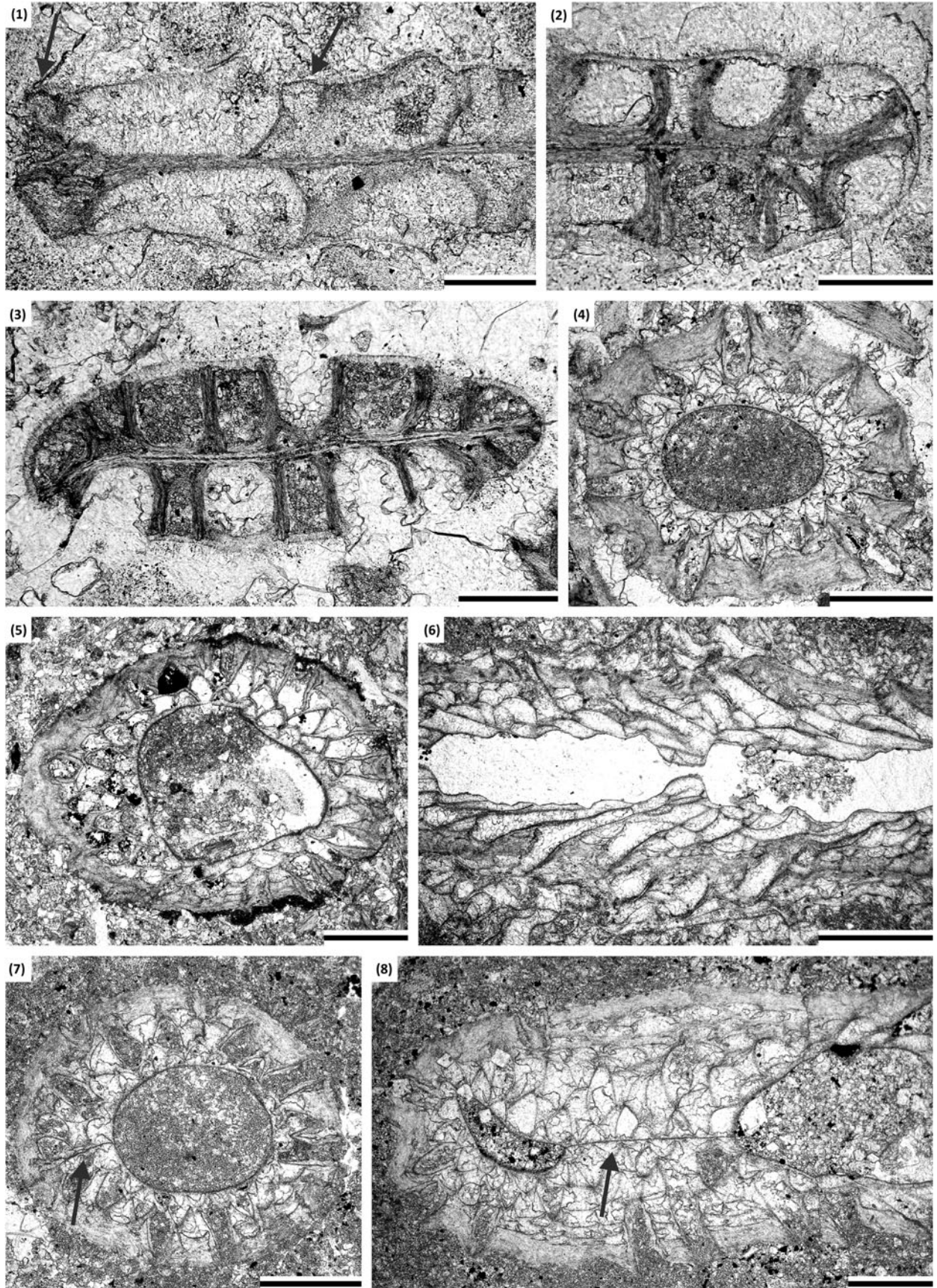
*Occurrence*.—Osgood Formation (Telychian-Sheinwoodian), Silurian; Indiana, USA. Reynales Formation, Hickory Corners Member, Aeronian, Llandoverly, lower Silurian; Hickory Corners, New York, USA.

*Description*.—Branched colonies, branch diameter 2.20–6.10 mm. Exozone 0.27–1.10 mm wide, endozone 1.56–3.94 mm wide, distinct. Secondary overgrowths uncommon. Autozoecia long in the endozone, having larger diameters than in exozone, polygonal in transverse section of endozone, bending gently and intersecting branch surface at angles of 57°–62°. Autozoecial apertures rounded polygonal. Autozoecial diaphragms rare to absent in the endozone, concentrated mostly in transitional region between endo- and exozone, common to abundant in outer exozone. Cystiphagms occurring throughout the exozone, occupying about half of the autozoecial diameter. Mesozooecia rare, locally common, small, short, restricted to the outermost part of exozone, polygonal, containing no diaphragms. Acanthostyles rare to common, large, prominent, restricted to exozone, with narrow hyaline cores and wide laminated sheaths. Autozoecial walls straight, locally weakly undulating, displaying granular microstructure, 0.002–0.003 mm thick in endozone; finely laminated with distinct medial lining, 0.05–0.11 mm thick in exozone.

*Etymology*.—The species name refers to the Niagara River, in which vicinity the type locality is situated.

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**Figure 5.** (1–9). *Diploclema argutum* Bassler, 1906: (1, 2) longitudinal section of a basal part of a colony encrusting *Moyerella* n. sp., SMF 23.327; (3) basal part of a colony encrusting sediment, SMF 23.344; (4) oblique longitudinal section showing autozoecia and interzoecial space (arrows), SMF 23.342; (5, 6) shallow tangential section showing autozoecial apertures, internal walls, and interzoecial space (arrows); (5) SMF 23.353; (6) SMF 23.354; (7) oblique section showing internal walls and interzoecial space, SMF 23.323; (8, 9) shallow tangential section showing autozoecial apertures, internal walls, and interzoecial space (arrows), SMF 23.323. (1) Scale bar = 1 mm; (2, 3, 7, 8) scale bars = 0.5 mm; (4–6, 9) scale bars = 0.2 mm.





**Table 1.** Descriptive statistics of *Diploclema argutum* Bassler, 1906. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Branch width	10	0.38	0.061	16.16	0.30	0.46
Autozooeical width	20	0.16	0.018	11.20	0.13	0.20
Autozooeical length	7	0.48	0.062	12.96	0.40	0.60
Interior wall thickness	10	0.02	0.008	40.75	0.01	0.04
Aperture width	7	0.08	0.013	15.66	0.07	0.10

*Remarks.*—*Homotrypa niagarensis* n. sp. differs from *H. anticostiensis* Bassler, 1928 from the Upper Ordovician of Canada in larger autozooeical apertures (aperture width 0.13–0.20 mm versus 0.09–0.14 mm in *H. anticostiensis*) and in absence of maculae consisting of mesozooeicia. *Homotrypa niagarensis* n. sp. differs from *H. ramulosa* Bassler, 1903 from the Upper Ordovician of Indiana, USA, in larger acanthostyles and larger autozooeical apertures (aperture width 0.13–0.20 mm versus 0.10–0.13 mm in *H. ramulosa*).

Family Halloporidae Bassler, 1911  
Genus *Hallopora* Bassler, 1911

*Type species.*—*Callopora elegantula* Hall, 1852, lower Silurian (Niagaran); USA.

*Hallopora* aff. *H. elegantula* (Hall, 1852)  
Figure 9.2–9.6; Table 5

*Occurrence.*—Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA.

*Description.*—Ramosely colonies, branch diameter 2.60–6.60 mm. Exozone distinct, 0.45–1.40 mm wide, endozone 1.70–4.62 mm wide. Secondary overgrowths common, 0.8–1.3 mm thick. Autozooeicia long, growing parallel to branch axis for a long distance in endozone, having rounded polygonal shape in transverse section in endozone, in exozone bending sharply and intersecting branch surface at angles of 60°–90°. Autozooeical apertures rounded to oval. Autozooeical diaphragms thin, planar, rare to common in endozone, becoming common in exozone, planar, rarely inclined, developed as extension of wall cortex. Cap-like apparatus and mural spines absent. Mesozooeicia arising in endozone, polygonal in cross section, often separating autozooeicia completely from each other in the exozone, 5–13 surrounding each autozooeical aperture. Mesozooeical diaphragms planar, densely spaced. Autozooeical walls indistinctly laminated, 0.005–0.010 mm thick in endozone; displaying distinct reverse V-shaped structure with dark autozooeical border,

having well-developed wall cortex continued in diaphragms, 0.04–0.08 mm thick in exozone. Maculae consisting of macrozooeicia, 1.1–1.5 mm in diameter.

*Materials.*—SMF 23.503–SMF 23.534.

*Remarks.*—The present species is very similar to *H. elegantula* (Hall, 1852). However, it differs in having smaller autozooeical apertures (0.27–0.32 mm versus 0.30–0.35 mm in *H. elegantula*) and smaller distances between aperture centers (0.30–0.47 mm versus 0.45–0.65 mm in *H. elegantula*). *Hallopora* aff. *H. elegantula* (Hall, 1852) is very similar to *H. baltica* (Hennig, 1908) from the Silurian (Wenlock) of Gotland but differs in rare diaphragms in endozone and in closely spaced autozooeical apertures (approximately six apertures per 2 mm distance versus four in *H. baltica*).

The present species is also similar to the species *Hallopora magnopora* Foerste, 1887, which was originally described from the lower Silurian of Ohio and redescribed by Bassler (1906) from the Rochester Shale (Wenlock) of New York. Both authors referred to “zooeicia larger than is usual in species of this genus” (Foerste, 1887, p. 173; Bassler, 1906, p. 43). However, they did not provide exact measurements of the apertures. Foerste (1887, p. 173) provided spacing of apertures as eight in 3 mm, and Bassler (1906, p. 43) gave the value of four apertures per 2 mm. Aperture width measured from the illustrations by Bassler (1906, pl. 15, fig. 1) are estimated as 0.35–0.40 mm, and these values are larger than in the present species (0.27–0.32 mm). Furthermore, according to illustrations in Bassler (1906, pl. 15, figs. 1–8), *H. magnopora* has fewer mesozooeicia than the present species.

Family Heterotrypidae Ulrich, 1890  
Genus *Leioclema* Ulrich, 1882  
[= *Lioclema* Ulrich, 1882]

*Type species.*—*Callopora punctata* Hall, 1858. Mississippian; Iowa, USA.

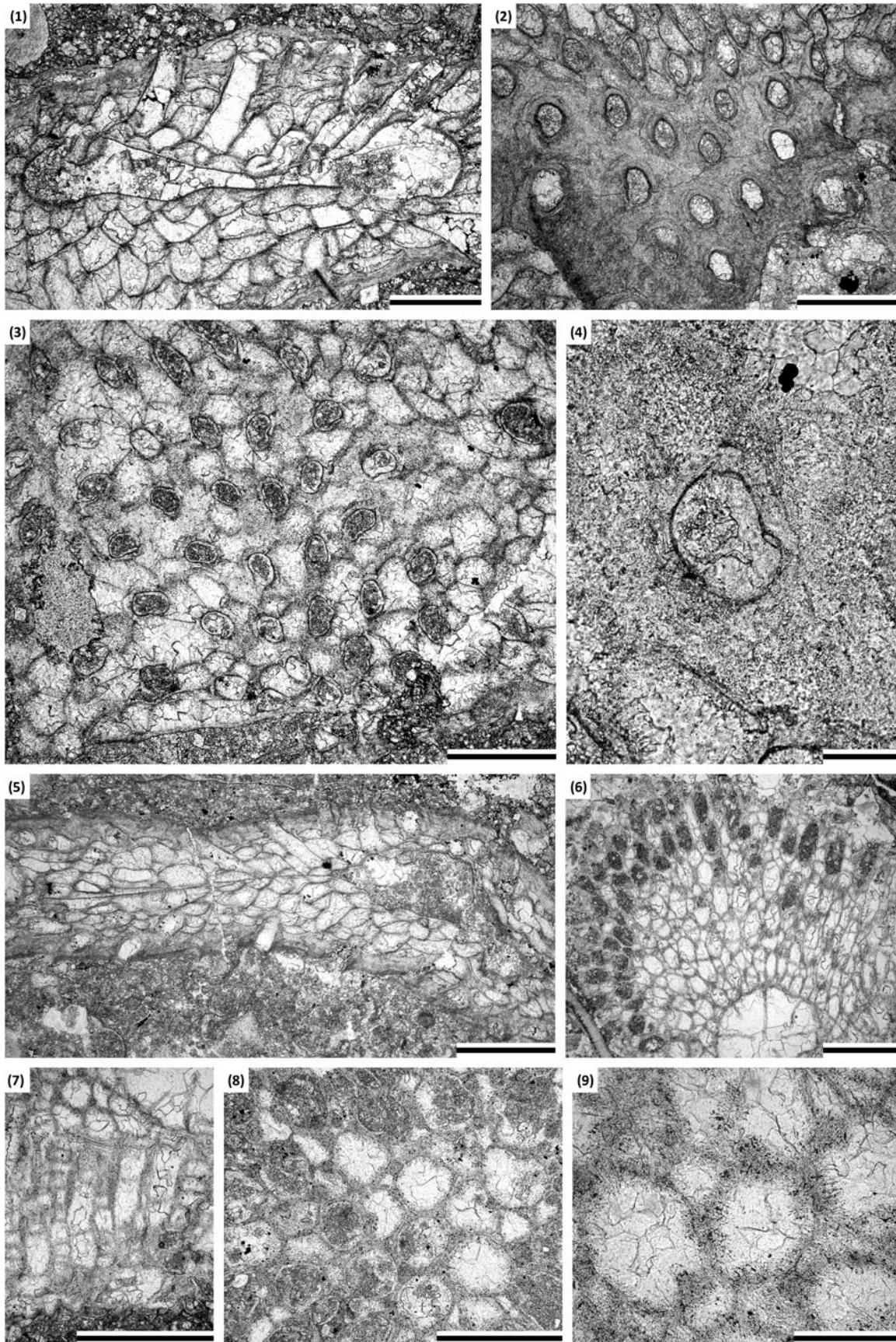
*Leioclema adsuetum* new species  
Figures 9.7, 10.1–10.5; Table 6

*Holotype.*—SMF 23.535.

*Paratypes.*—SMF 23.536–SMF 23.558.

*Diagnosis.*—Ramosely branched colonies with narrow distinct exozones; secondary overgrowth occurring; autozooeicia long, bending at mediate angles in exozone, having rounded polygonal apertures; basal diaphragms rare to absent in exozone; mesozooeicia abundant, four to nine surrounding

**Figure 6.** (1–3) *Diploclema argutum* Bassler, 1906: (1) longitudinal section showing autozooeical chambers, external walls, internal walls, and protruded interzooeical space in proximal part of autozooeicia (arrows), SMF 23.333; (2) branch transverse section showing autozooeical chambers, external and internal walls, SMF 23.351; (3) branch transverse section showing autozooeical chambers, external and internal walls, SMF 23.358. (4–8) *Cheilotrypa* aff. *C. variolata* (Hall, 1876): (4) transverse section of a hollow cylindrical colony, SMF 23.436; (5) transverse section of a hollow cylindrical colony, SMF 23.402; (6) longitudinal section of a tubular colony, SMF 23.423; (7, 8) transverse section of a hollow cylindrical colony with sheets budding “base to base” (arrow), SMF 23.428. (1–3) Scale bars = 0.2 mm; (4, 5, 7, 8) scale bars = 0.5 mm; (6) scale bar = 1 mm.



**Table 2.** Descriptive statistics of *Cheilotrypa* aff. *C. variolata* (Hall, 1876). N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value.

	N	X	SD	CV	Min	Max
Colony thickness, mm	8	0.56	0.264	47.51	0.26	1.10
Aperture width, mm	20	0.11	0.013	11.52	0.09	0.15
Aperture spacing, mm	20	0.28	0.030	10.51	0.24	0.36
Vesicle diameter, mm	20	0.11	0.027	24.44	0.07	0.16
Vesicles per aperture	20	6.0	0.945	15.87	5.0	8.0

each autozoecial aperture; acanthostyles moderately large and abundant, two to five surrounding each autozoecial aperture; autozoecial walls laminated, moderate in thickness.

**Occurrence.**—Osgood Formation (Telychian-Sheinwoodian), Silurian; Indiana, USA. Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA.

**Description.**—Ramosely branched colonies, branch diameter 2.55–5.88 mm. Exozone distinct, 0.34–0.72 mm wide, endozone 1.48–4.44 mm wide. Secondary overgrowth common, 0.3–0.4 mm thick. Autozoecia long, growing parallel to branch axis for a long distance in endozone, having larger diameters than in exozone, polygonal in transverse section of endozone, in exozone bending sharply and intersecting branch surface at angles of 80°–90°, having polygonal shape in cross section in endozone. Autozoecial apertures polygonal. Autozoecial diaphragms rare to absent, thin, straight or slightly deflected proximally. Mesozoecia abundant, four to nine surrounding each aperture, polygonal in cross section, strongly beaded, containing abundant planar diaphragms. Acanthostyles moderately large, abundant, two to five surrounding each aperture, originating in the distal part of exozone, often indenting autozoecia, having distinct calcite cores and dark laminated sheaths. Walls granular, 0.005–0.008 mm thick in endozones; distinctly laminated, 0.040–0.045 mm thick in the exozone.

**Etymology.**—The species name refers to the average appearance of this species (from Latin *adsuetus* meaning normal, usual).

**Remarks.**—*Leioclema adsuetum* n. sp. differs from *L. ramulosum* (Bassler, 1906) from the Rochester Shale of New York in more abundant acanthostyles surrounding autozoecial apertures (2–5 versus 1–2 in *L. ramulosum*) and in having abundant diaphragms in mesozoecia. *Leioclema adsuetum* n. sp. differs from *L. densiporum* Owen, 1965 from the lower Silurian of England in having less abundant acanthostyles surrounding autozoecial apertures (2–5 versus 8–12 in *L. densiporum*).

Order Cryptostomata Vine, 1884

Suborder Rhabdomesina Astrova and Morozova, 1956

**Table 3.** Descriptive statistics of *Hennigopora apta* Perry and Hattin, 1960. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Aperture width	20	0.19	0.018	9.52	0.17	0.25
Aperture spacing	20	0.30	0.040	13.17	0.23	0.40
Acanthostyle diameter	20	0.054	0.008	15.03	0.040	0.065
Vesicle diameter	20	0.10	0.027	27.22	0.05	0.15
Vesicle spacing	15	0.10	0.030	29.10	0.05	0.16

Family Arthrostylidae Ulrich, 1882  
Genus *Moyerella* Nekhoroshev, 1956

**Type species.**—*Moyerella stellata* Nekhoroshev, 1956. Llandovery, lower Silurian; Russia (Siberia).

**Diagnosis (emended).**—Thin ramose colonies, usually articulated. Axial region consisting of well-defined median axis or planar budding surface. Autozoecia shortened tubular, triangular in cross section in endozone, inflated at their bases, diverging from median axis in spiral pattern, bending abruptly in exozones. Diaphragms rare or absent. Hemisepta absent. Autozoecial apertures oval or rounded, arranged regularly in diagonal rows, commonly having peristomes. Zoecial boundaries well defined, narrow. Single large four-sided acanthostyle between longitudinally successive autozoecial apertures present. Paired heterozoecia (tectitooecia after Kopajevich, 1975) present between successive autozoecial apertures. Tubules often present in peristomes.

**Occurrence.**—Upper Ordovician of Argentina and Estonia; lower Silurian (Llandovery) of Siberia and Estonia and United States.

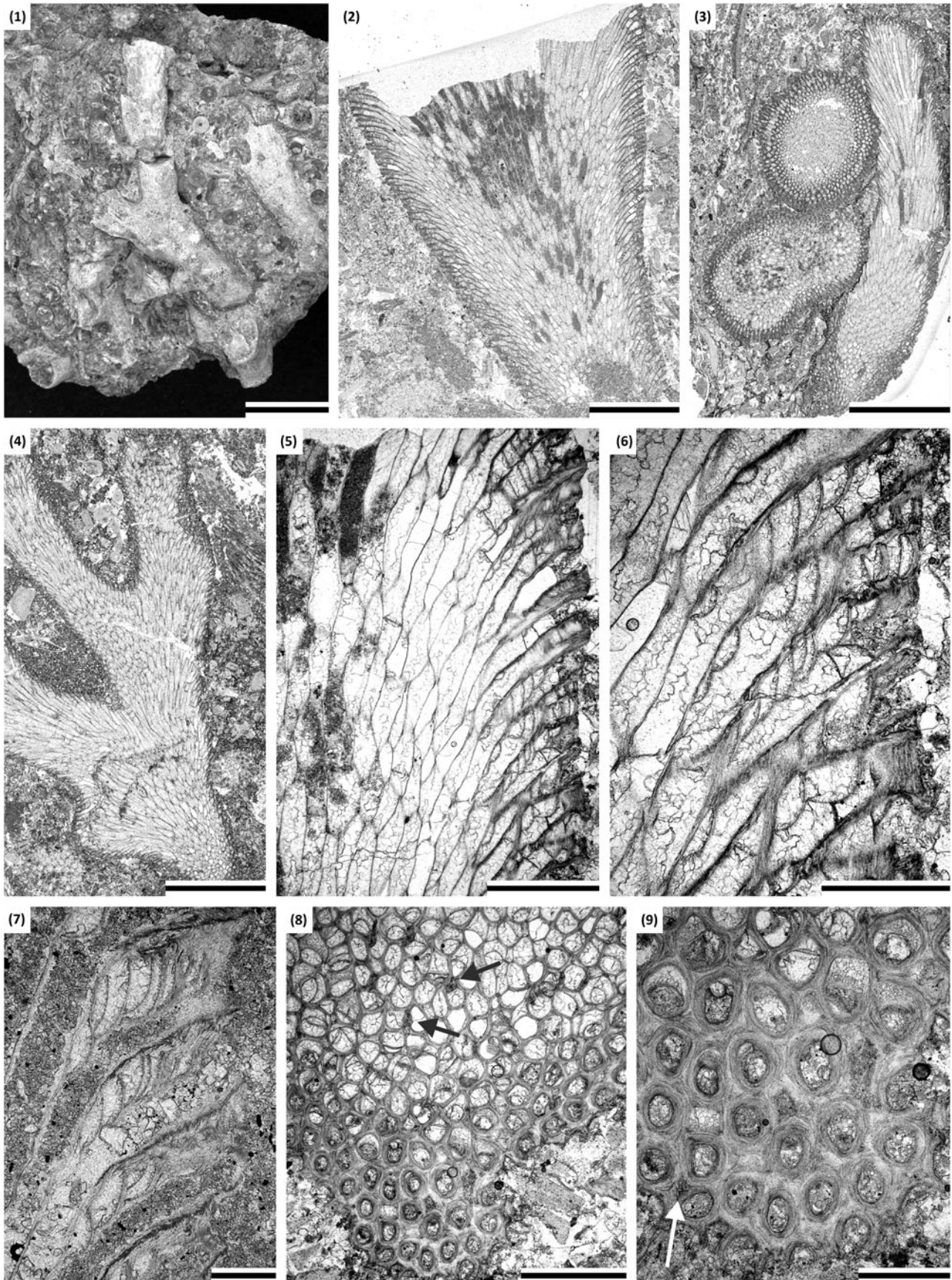
**Remarks.**—*Moyerella* Nekhoroshev, 1956 is similar to *Nematotrypa* Bassler, 1911 in autozoecial shape and presence of acanthostyles and heterozoecia with diaphragms (tectitooecia). However, *Moyerella* differs from *Nematotrypa* in the regular arrangement of paired heterozoecia between autozoecia and absence of hemiphragms.

Blake (1983, p. 561) mentioned paurostyles in the diagnosis of *Moyerella*. However, this kind of style is characterized by simple hyaline cores, whereas *Moyerella* has tubules that are deflections of the laminated skeletal material. Nekhoroshev (1956, p. 47, pl. 11, fig. 3a) mentioned the presence of granules on ridges between apertures that are indeed tubules.

*Moyerella parva* new species  
Figures 10.6, 11.1–11.10; Table 7

**Holotype.**—SMF 23.559.

**Figure 7.** (1–5) *Cheilotrypa* aff. *C. variolata* (Hall, 1876): (1) oblique section of a hollow cylindrical colony, SMF 23.423; (2) shallow tangential section showing autozoecial apertures with lunaria, SMF 23.379; (3) deep tangential section showing autozoecial apertures with lunaria and vesicles, SMF 23.423; (4) tangential section of autozoecial aperture, SMF 23.423; (5) oblique section of a hollow cylindrical colony, SMF 23.425. (6–9) *Hennigopora apta* Perry and Hattin, 1960: (6) oblique section of a colony, SMF 23.466; (7) longitudinal section showing autozoecia and vesicles, SMF 23.466; (8, 9) tangential section showing autozoecia, acanthostyles, and vesicles; (8) SMF 23.462; (9) SMF 23.465. (1–3, 8) Scale bars = 0.5 mm; (4) scale bar = 0.1 mm; (5–7) scale bars = 1 mm; (9) scale bar = 0.2 mm.



*Paratypes*.—SMF 23.560–SMF 23.633.

*Diagnosis*.—Articulated colonies consisting of segments with pointed bases and widened proximal parts, apparently flexibly connected; segments cylindrical with narrow exozones; autozoecia moderately long, budding at low angles from the medial axis, triangular in endozone, bending abruptly in exozone; autozoecial diaphragms rare; paired heterozoecia regularly arranged between longitudinally successive autozoecial apertures; single acanthostyle between heterozoecia present; low ridges between apertures containing tubules.

*Occurrence*.—Osgood Formation (Telychian-Sheinwoodian), Silurian; Indiana, USA. Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA.

*Description*.—Articulated colonies consisting of segments with pointed bases and widened proximal parts, apparently flexibly connected; segments cylindrical, 0.4–1.0 mm in diameter, with 0.1–0.3 mm wide exozones and 0.2–0.4 mm wide endozones. Branching unknown. Medial axis well developed. Autozoecia moderately long, inclined at angles of 16°–25° to the medial axis, having triangular shape in cross section in endozone, bending abruptly in exozone and intersecting the branch surface at angles of 49°–68°. Autozoecial diaphragms rare. Autozoecial apertures oval, arranged in distinct diagonal rows. Paired heterozoecia (tectitooecia) arranged regularly between longitudinally successive autozoecial apertures, short, forming conical tubes, having triangular to pentagonal shape in cross section, 0.025–0.045 mm in width. Diaphragms in heterozoecia not observed. Autozoecial walls hyaline, 0.008–0.010 mm thick in endozone; reverse V-shaped laminated, 0.045–0.063 mm thick in exozone. Single acanthostyle between longitudinally successive autozoecial apertures present, 0.020–0.025 mm in diameter. Low ridges on the colony surface between autozoecial apertures producing four-sided pattern at junctions where acanthostyles are present. Dark tubules on ridges, 0.008–0.010 mm in diameter.

*Etymology*.—The species name refers to the small size of this species (from Latin *parva* meaning small).

*Remarks*.—*Moyerella parva* n. sp. is similar to *M. stellata* Nekhoroshev, 1956 from the lower Silurian (Llandovery) of Siberia but differs in possessing smaller diameter segments (0.40–1.00 mm versus 0.45–1.65 mm in *M. stellata*) and smaller autozoecial apertures (aperture width 0.06–0.09 mm versus 0.09–0.10 mm in *M. stellata*). Moreover, tectitooecia of *M. stellata* have two to four diaphragms, whereas no

diaphragms were observed in tectitooecia of *M. parva* n. sp. Species *Helopora formosa* Billings, 1866 from the Jupiter Formation (Llandovery) of the Anticosti Island is superficially similar to *M. parva* n. sp. in the shape of segments; however, no further details are known about this species.

Suborder Ptilodictyina Astrova and Morozova, 1956

Family Ptilodictyidae Zittel, 1880

Genus *Phaenopora* Hall, 1851

[ = *Fimbriopora* Astrova, 1965 ]

*Type species*.—*Phaenopora explanata* Hall, 1852. Silurian; Canada.

*Phaenopora multifida* (Hall, 1883)

Figures 11.11–11.14, 12.1–12.3; Table 8

1883 *Stictopora multifida* Hall, p. 268, pl. 14, fig. 4.

1887 *Phaenopora multifida* (Hall, 1883); Foerste, p. 160.

1888 *Phaenopora multifida* (Hall, 1883); Foerste, pl. 16, fig. 3.

1893 *Phaenopora multifida* (Hall, 1883); Foerste, p. 599, pl. 29, fig 3.

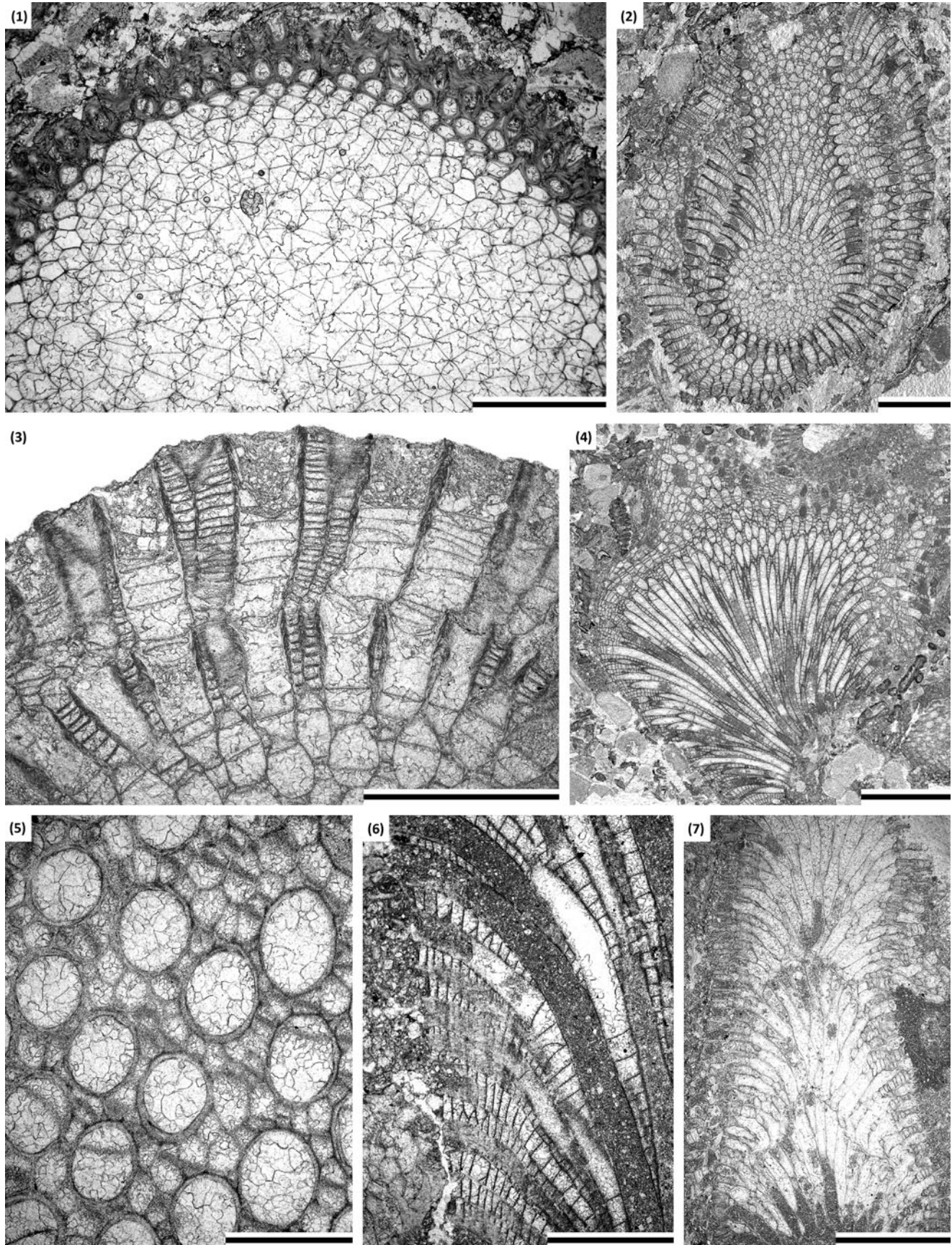
*Occurrence*.—Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA. Clinton Group, Silurian; Ohio, USA.

*Description*.—Broad and flattened bifoliate branches, 1.0–4.5 mm wide and 0.35–1.15 mm thick. Mesotheca zigzag folded in transverse section, three layered, 0.013–0.020 mm thick. Autozoecia short, rectangular in deep tangential section. Autozoecial apertures oval to rectangular with rounded corners, arranged in 8–13 rows. Superior and observer hemisepta present, long. Metazooecia small, shallow, having triangular to pentagonal apertures, arranged regularly in pairs between apertures. Autozoecial walls finely laminated, 0.010–0.013 mm thick in endozones; laminated, fused, thick in exozones. Wide longitudinal crests on colony surface present.

*Materials*.—SMF 23.634–SMF 23.689.

*Remarks*.—*Phaenopora multifida* (Hall, 1883) differs from *P. markhensis* Nekhoroshev, 1961 from the lower Silurian (Llandovery) of Siberia in its smaller autozoecial apertures (aperture width 0.08–0.10 mm versus 0.10–0.13 mm in *P. markhensis*). *Phaenopora multifida* differs from *P. contracta* Astrova, 1959 from the Wenlock of Tuva (Russia) in thinner and narrower colonies and larger autozoecial apertures (aperture width 0.08–0.10 mm versus 0.06–0.07 mm in *P. contracta*).

**Figure 8.** (1–9) *Homotrypa niagarensis* n. sp.: (1) colony in rock, holotype SMF 23.474; (2) branch oblique section, holotype SMF 23.470; (3) rock thin section with transverse and oblique sections of branches, holotype SMF 23.472; (4) branch oblique section, paratype SMF 23.488; (5, 6) longitudinal section of exozone showing autozoecia with diaphragms and cystiphagms, holotype SMF 23.470; (7) longitudinal section of outer exozone showing autozoecial chambers with series of cystiphagms, paratype SMF 23.483; (8, 9) tangential section showing autozoecial apertures and acanthostyles (arrow), holotype SMF 23.473. (1) Scale bar = 10 mm; (2) scale bar = 3 mm; (3, 4) scale bars = 5 mm; (5, 8) scale bars = 1 mm; (6, 9) scale bars = 0.5 mm; (7) scale bar = 0.2 mm.



**Table 4.** Descriptive statistics of *Homotrypa niagarensis* n. sp. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Branch width	10	3.73	1.288	34.59	2.20	6.10
Exozone width	10	0.54	0.249	46.39	0.27	1.10
Endozone width	10	2.65	0.836	31.52	1.56	3.94
Aperture width	30	0.13	0.023	17.40	0.10	0.20
Aperture spacing	30	0.23	0.030	12.88	0.16	0.30
Mesozoecia width	30	0.06	0.010	18.34	0.03	0.07
Acanthostyle diameter	25	0.09	0.017	18.57	0.06	0.12
Exozonal wall thickness	30	0.07	0.019	26.84	0.05	0.11
Cystiphragm spacing	30	0.07	0.025	35.24	0.03	0.13

Genus *Ptilodictya* Lonsdale, 1839

*Type species.*—*Flustra lanceolata* Goldfuss, 1829. Lower Silurian (Wenlock); Great Britain.

*Ptilodictya sulcata* Billings, 1866  
Figure 12.4–12.8; Table 9

1866 *Ptilodictya sulcata* Billings, p. 35.

1911 *Ptilodictya gladiola*; Bassler, p. 114, fig. 43.

1928 *Ptilodictya sulcata*; Bassler, p. 162, pl. 10, figs. 9–11.

1962 *Ptilodictya gladiola*; Kieppura, p. 405, pl. 9, fig. 1.

*Lectotype.*—GSC 2501. Jupiter Formation (Aeronian–Telychian), Llandoverly, lower Silurian; Anticosti, Canada.

*Occurrence.*—Reynales Formation, Hickory Corners Member, Aeronian, Llandoverly, lower Silurian; Hickory Corners, New York, USA. Upper Ordovician to lower Silurian; Anticosti, Canada. Porkuni Stage (Hirnantian); Estonia.

*Description.*—Bifoliate branched colony with pointed base, 0.72–1.35 mm wide and 0.49–0.52 mm thick in its distal part. Mesotheca zigzag folded, three layered, 0.015–0.020 mm thick; median rods lacking. Autozoecia moderately long, rectangular in deep tangential section. Autozoecial apertures rectangular with rounded corners, arranged in eight or nine rows. Short hemisepta present. Occasional thin diaphragms occurring. Heterozoecia and styles absent. Autozoecial walls in endozone laminated with dark dividing layer, 0.018–0.023 mm thick; in exozone coarsely laminated. Maculae not observed.

*Materials.*—SMF 23.690–SMF 23.694.

*Remarks.*—The present material is similar to *Ptilodictya sulcata* Billings, 1866. Ross (1960b) redescribed and depicted the lectotype as well as extensive additional material from the Upper Ordovician and lower Silurian of Anticosti. The present material differs in having narrower branches (branch width 0.72–1.35 mm versus 1.2–3.3 mm in the Canadian material).

**Table 5.** Descriptive statistics of *Hallopora* aff. *H. elegantula* (Hall, 1852). N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value.

	N	X	SD	CV	Min	Max
Branch width, mm	7	4.92	1.722	35.00	2.60	6.60
Exozone width, mm	7	0.87	0.338	38.77	0.45	1.40
Endozone width, mm	7	3.18	1.249	39.34	1.70	4.62
Aperture width, mm	30	0.27	0.020	7.53	0.23	0.32
Aperture spacing, mm	30	0.37	0.043	11.84	0.30	0.47
Macrozoecia width, mm	20	0.34	0.020	5.92	0.30	0.38
Mesozoecia width, mm	30	0.10	0.034	33.46	0.03	0.17
Mesozoecia per aperture	20	8.9	1.981	22.38	5.0	13.0
Mesozoecial diaphragm spacing, mm	30	0.08	0.021	25.05	0.05	0.14

However, internal characteristics and aperture dimensions are identical; therefore, we identify the material from the Reynales Formation as *P. sulcata*.

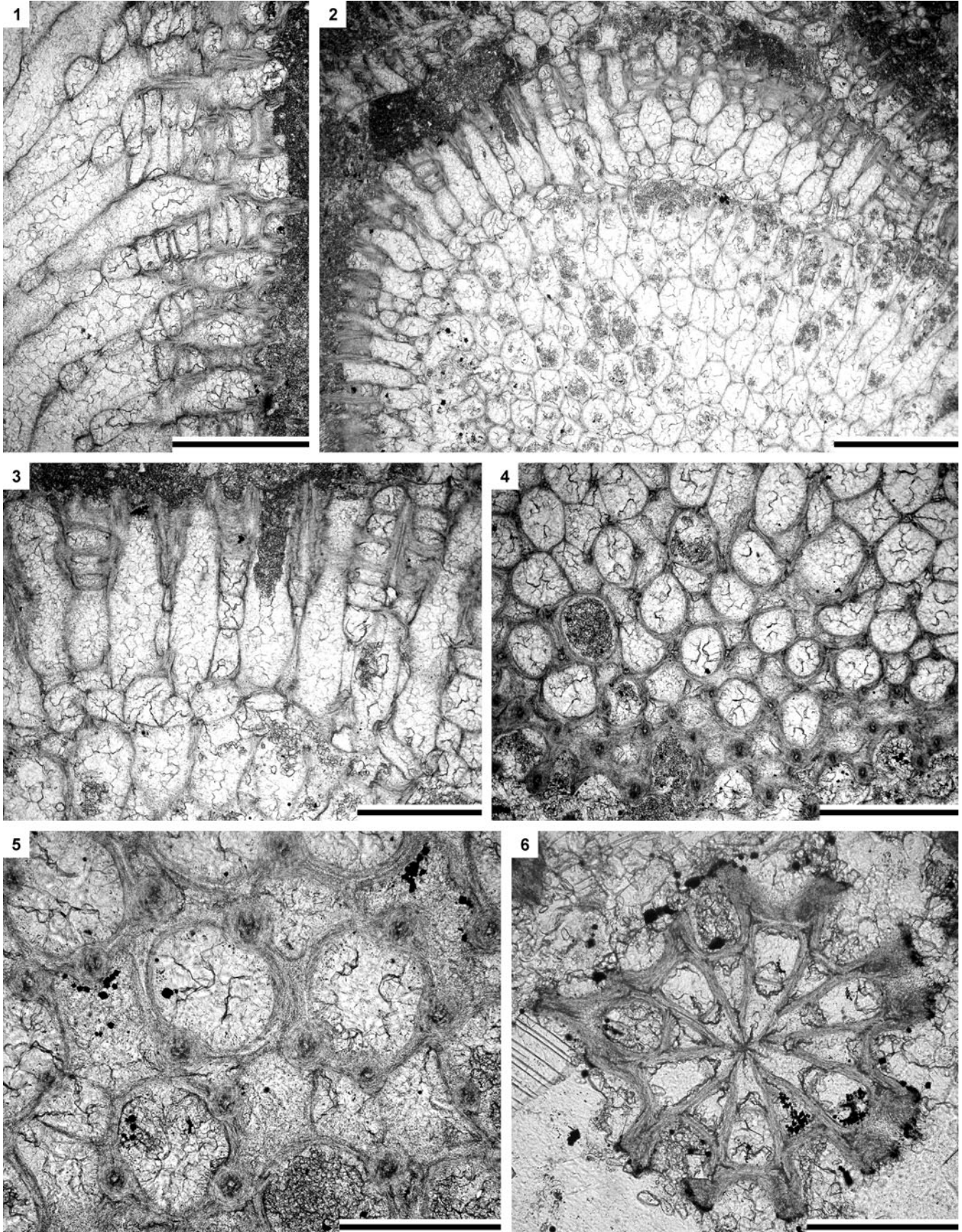
Two more species of *Ptilodictya* were established from the Ordovician–Silurian deposits of North America that are quite close in their characteristics to *P. sulcata*: *P. ensiformis* (Hall, 1852) and *P. gladiola* Billings, 1866. Ross (1960b, p. 1068) studied the holotype of *P. gladiola* and supposed it being a synonym of *P. ensiformis*. This holotype was in bad condition, so Ross (1960b) could not find any morphological features for the species determination. She noted also that the majority of the material referred by Bassler (1928) to *P. gladiola* is in fact *Stictoporella (Phaenopora) excellens* (Billings, 1866).

A general problem in the study of the lower Paleozoic cryptostomes is that many genera possess similar growth forms and dimensions but differ in their internal morphology. Inadequate descriptions and illustrations of original publications produce additional difficulty in discrimination of species. In some cases, study of the type material (or adequate material from type areas) has been possible. Material found at the National Museum of Natural History, Washington, D.C., contained three sets of specimens referred to *Ptilodictya ensiformis* (USNM PAL 35752, USNM PAL 69042, and USNM PAL 71560) and one set referred to *P. gladiola* (USNM PAL 57210). The latter material was referred by Bassler (1911) from the Upper Ordovician of Estonia. This specimen is accompanied by a tangential thin section that reveals characteristic features of *Ptilodictya*. Dimensions and internal morphology of this specimen are close to those of the material from the Reynales Formation (width of autozoecial apertures 0.09–0.10 mm versus 0.07–0.08 mm in *P. sulcata*; spacing of apertures alongside the branch 0.21–0.29 mm versus 0.22–0.31 mm *P. sulcata*).

Study of photographs of the hypotype USNM PAL 69042 at the National Museum of Natural History, Washington, D.C., referred by Bassler (1928) to *Ptilodictya ensiformis* (Ordovician–Silurian of Anticosti), revealed the presence of two metazoecia between successive autozoecial apertures; therefore, this material is regarded as being *Phaenopora*. This species developed similar growth form with a pointed base, and it has dimensions similar to *Ptilodictya sulcata* and *P. ensiformis*.

**Figure 9.** (1) *Homotrypa niagarensis* n. sp.: part of branch transverse section showing thin-walled autozoecia in endozone and thick-walled autozoecia in exozone, holotype SMF 23.471. (2–6) *Hallopora* aff. *H. elegantula* (Hall, 1852): (2) oblique section of branch with secondary overgrowth, SMF 23.504; (3) longitudinal section of exozone showing secondary overgrowth, autozoecia with diaphragms, and mesozoecia, SMF 23.504; (4) oblique section of branch showing long zooecia in endozone, SMF 23.531; (5) tangential section showing autozoecial apertures and mesozoecia, SMF 23.530; (6) longitudinal section showing autozoecia and mesozoecia, SMF 23.532. (7) *Leioclema adsuetum* n. sp.: longitudinal section of a branched colony with secondary overgrowth, paratype SMF 23.553. (1, 3, 6) Scale bars = 1 mm; (2, 7) scale bars = 2 mm; (4) scale bar = 3 mm; (5) scale bar = 0.5 mm.





**Table 6.** Descriptive statistics of *Leioclema adsuetum* n. sp. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value.

	N	X	SD	CV	Min	Max
Branch width, mm	8	3.38	1.142	33.79	2.25	5.88
Exozone width, mm	8	0.53	0.134	25.14	0.34	0.72
Endozone width, mm	8	2.32	0.943	40.72	1.48	4.44
Aperture width, mm	30	0.14	0.020	13.96	0.11	0.19
Aperture spacing, mm	30	0.23	0.031	13.47	0.18	0.31
Acanthostyle diameter, mm	30	0.05	0.009	16.77	0.04	0.07
Mesozoocelia width, mm	30	0.09	0.020	23.68	0.05	0.14
Acanthostyles per aperture	30	3.6	0.765	21.05	2.0	5.0
Mesozoocelia per aperture	30	6.2	1.234	20.01	4.0	9.0
Mesozoocelial diaphragm spacing, mm	30	0.10	0.031	30.47	0.05	0.15

The hypotype USNM PAL 35752 for *Ptilodictya ensiformis* (Hall, 1852), referred by Bassler (1906) from the Rochester Shale (Wenlock) of New York, shows typical features of *Ptilodictya*, but this material differs from the present species in larger autozoocelial apertures (width of autozoocelial apertures 0.13–0.17 mm versus 0.07–0.08 mm in *P. sulcata*) and wider spacing of apertures alongside the branch (0.43–0.53 mm versus 0.22–0.31 mm *P. sulcata*).

The specimen USNM PAL 71560, referred to by Butts (1926) as *Ptilodictya ensiformis* (Hall, 1852) from the Paleozoic of Alabama, is indeed *Phaenopora*. This specimen has wide branches and two metazoocelia between autozoocelial apertures.

The species described by Kiepara (1962) as *Ptilodictya gladiola* Billings, 1866 from erratic boulders of Poland shows similarities with the present material. It has branches of 0.8 mm width and apertures of ~0.08 mm width (measured from fig. 1 on pl. 9 in Kiepara, 1962), which are close to present species. Internal morphology of the Polish material is unknown.

*Ptilodictya sulcata* is similar to *P. exiliformis* Suttner and Ernst, 2007 from the Upper Ordovician of India (Pin Formation). The latter species has narrower branches (branch width 0.41–0.81 mm versus 0.72–1.35 mm in present material [up to ~3 mm given by other authors]). *Ptilodictya* differs from *P. exilis* Lavrentjeva in Gorjunova and Lavrentjeva, 1993 from the Upper Ordovician (Katian) of Estonia in having smaller autozoocelial apertures (aperture width 0.07–0.08 mm versus 0.10–0.13 mm in *P. exilis*).

Family Rhinidictyidae Ulrich, 1893  
Genus *Metadictya* Kopajevich, 1975

*Type species.*—*Hemipachydictya copiosa* Kopajevich, 1970. Ordovician–Silurian (Ashgill, Llandovery); Estonia.

*Metadictya* sp.

Figures 12.9, 12.10, 13.1–13.3; Table 10

*Occurrence.*—Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA.

*Description.*—Bifoliate branched colony, 1.5–2.5 mm wide and 0.60–0.65 mm thick. Mesotheca straight, three layered, 0.02–0.03 mm thick; median rods present. Autozoocelia moderately long, rectangular in deep tangential section. Autozoocelial apertures oval to rectangular with rounded corners, arranged in 9–13 rows. Hemisepta absent. Occasional thin diaphragms occur. Heterozoocelia and styles absent. Autozoocelial walls in endozone laminated with dark dividing layer, 0.015–0.020 mm thick; in exozone coarsely laminated, containing numerous spherules in autozoocelia walls, regularly arranged in transverse bands. Spherules 0.010–0.015 mm in diameter.

*Materials.*—SMF 23.695–SMF 23.704.

*Remarks.*—*Metadictya* sp. differs from *M. bifurcata* (Hall, 1883) from the lower Silurian (Llandovery) of Indiana in narrower and thinner branches (branch width 1.5–2.5 mm versus 4–5 mm in *M. bifurcata*; branch thickness 0.60–0.65 mm versus 2–3 mm in *M. bifurcata*). Foerste (1887, p. 163) described material *M. bifurcata* from the lower Silurian of Ohio, and he gave longitudinal spacing of 5–5.5 apertures per 2 mm distance, which would closely fit with the present species (distance between aperture centers 0.34–0.40 mm [measured] versus 0.37–0.40 mm [converted] in *M. bifurcata*). *Metadictya bifurcata* (Hall, 1883) has never been described and illustrated with use of thin sections from the type area, but this species was identified in some Ordovician localities in Europe (e.g., Bassler, 1911; Kiepara, 1962; Spjeldnaes, 1984). However, the relationship of these materials to *Metadictya bifurcata* (Hall, 1883) is questionable. *Metadictya* sp. differs from *M. striatipora* (Kopajevich, 1970) from the lower Silurian (Llandovery) of Estonia in narrower and thinner branches as well as in smaller autozoocelial apertures (aperture width 0.06–0.08 mm versus 0.11–0.28 mm in *M. striatipora*).

Genus *Trigonodictya* Ulrich, 1893  
[= *Astreptodictya* Karklins, 1969]

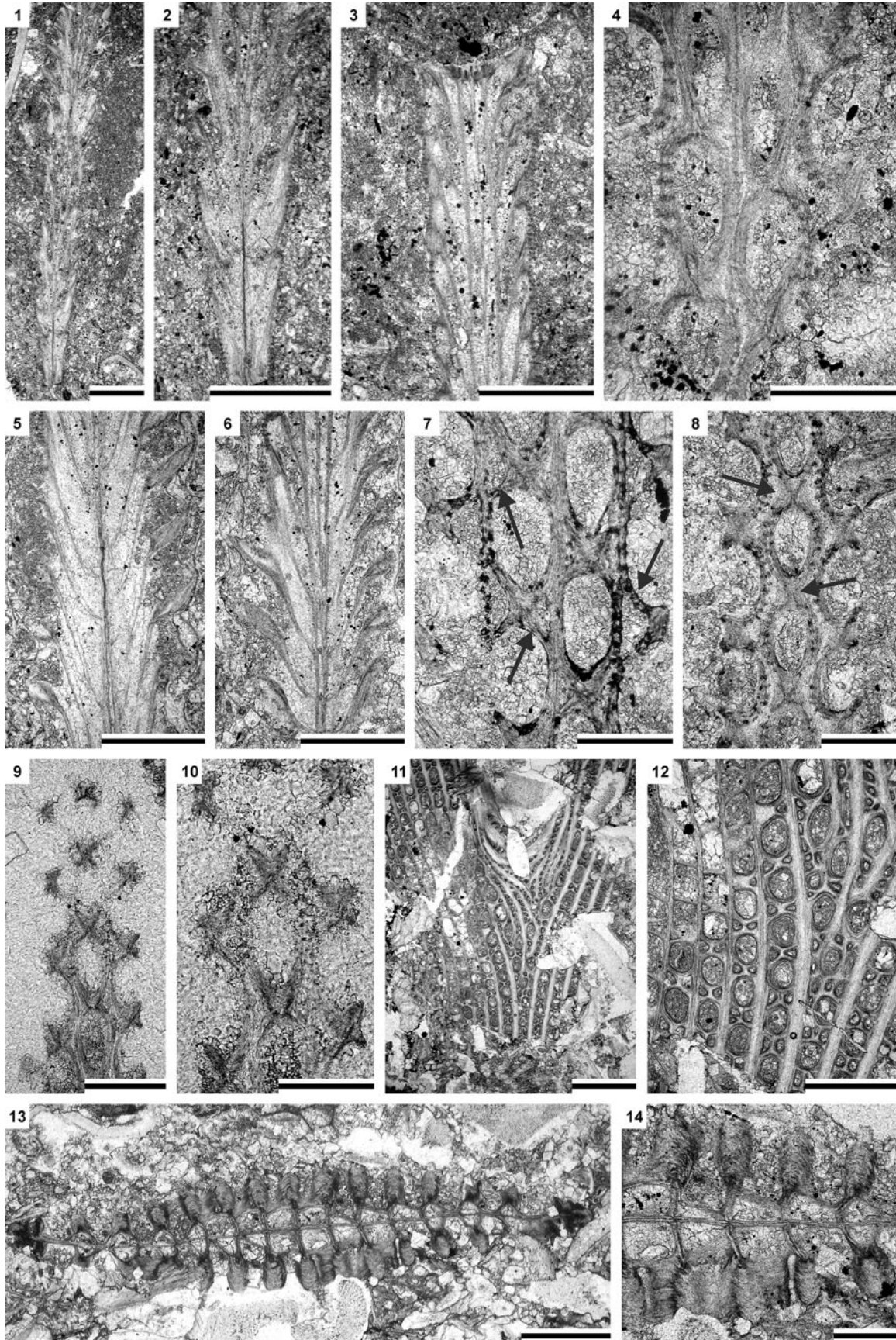
*Type species.*—*Pachydictya conciliatrix* Ulrich, 1886. Middle Ordovician (Decorah Shale); Minnesota, USA.

*Trigonodictya* sp.  
Figure 13.4–13.7; Table 11

*Occurrence.*—Reynales Formation, Hickory Corners Member, Aeronian, Llandovery, lower Silurian; Hickory Corners, New York, USA.

*Description.*—Branched bifoliate, dichotomous colonies. Branches flattened, with sharp edges, 3.5 mm wide and 0.97 mm thick. Mesotheca three layered, straight both in longitudinal and transverse sections, containing abundant median rods, 0.02–0.03 mm thick. Median rods densely spaced, 0.01–0.02

←  
**Figure 10.** (1–5) *Leioclema adsuetum* n. sp.: (1) longitudinal section of exozone showing autozoocelia, mesozoocelia, and acanthostyles, paratype SMF 23.553; (2, 3) oblique section showing autozoocelia in endozone and exozone and mesozoocelia, holotype SMF 23.535; (4, 5) tangential section showing autozoocelial apertures, mesozoocelia, and acanthostyles, holotype SMF 23.535. (6) *Moyerella parva* n. sp.: branch transverse section, paratype SMF 23.616. (1, 3, 4) Scale bars = 0.5 mm; (2) scale bar = 1 mm; (5, 6) scale bars = 0.2 mm.



**Table 7.** Descriptive statistics of *Moyerella parva* n. sp. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value.

	N	X	SD	CV	Min	Max
Branch width, mm	25	0.59	0.126	21.32	0.40	1.00
Exozone width, mm	25	0.15	0.045	29.29	0.10	0.30
Endozone width, mm	25	0.28	0.058	20.38	0.20	0.40
Aperture width, mm	20	0.08	0.009	11.61	0.06	0.09
Aperture spacing along branch, mm	20	0.28	0.022	7.79	0.23	0.32
Aperture spacing diagonally, mm	18	0.20	0.017	8.81	0.16	0.22
Autozoecial budding angle, endozone	12	19.3	2.934	15.17	16	25
Autozoecial budding angle, exozone	12	56.8	5.060	8.90	49	68

mm in diameter, continuous in dark zones separating longitudinal rows of autozoecia. Autozoecia regularly arranged in 10 alternating longitudinal rows, semicircular to trapezoid in transverse section in endozone, rectangular in deep tangential section, becoming oval on the colony surface. Autozoecial boundaries distinct, delineated laterally by continuous dark zones. Basal diaphragms rare or absent, straight. Extrazooecial skeletal deposits well developed, consisting of laminar and vesicular portions. Vesicular structures small, having flat to rounded roofs, rare to common in inner exozones. Laminar stereom with dark zones, longitudinally aligned, separating autozoecia in exozones. Monticules absent.

*Materials.*—SMF 23.705–SMF 23.709.

*Remarks.*—*Trigonodictya* sp. is most similar to *T. bifurcata instabilis* (Foerste, 1887) from Llandovery of Ohio and to *T. ambigua* (Ross, 1961) from the Upper Ordovician and lower Silurian of Canada. Foerste (1887, p. 164) provided the following measurements for his species: branch width 3–5 mm, branch thickness 1–2 mm, and six to seven apertures in 2 mm longitudinally. The latter value is larger than in the present species: six to seven versus four to five apertures per 2 mm. Ross (1961, p. 340) provided a comprehensive description of *T. ambigua* accompanied by a large set of measurements. *Trigonodictya* sp. differs from *T. ambigua* in larger autozoecial apertures (0.18–0.24 mm versus 0.08–0.20 mm in *T. ambigua*). Furthermore, *T. ambigua* possesses few and small vesicular structures in endozone, which are larger and more abundant in the present species.

Order Fenestrata Elias and Condra, 1957  
 Suborder Phylloporinina Lavrentjeva, 1979  
 Family Enalloporidae Lavrentjeva, 1985  
 Genus *Chasmatopora* Eichwald, 1855

*Type species.*—*Retepora tenella* Eichwald, 1855. Upper Ordovician, Ashgill; Vormsi, Estonia.

*Chasmatopora foerstei* McKinney and Wyse Jackson, 2010a  
 Figures 13.8, 14.1–14.3; Table 12

- non *Retepora angulata* Hall, p. 49, pl. 19, fig. 3a–h.  
 1852  
 ?1875 *Retepora angulata*?; Hall and Whitfield, p. 111, pl. 5, figs. 2–4.  
 1887 *Phylloporina angulata* (Hall, 1852); Foerste, p. 151, pl. 15, figs. 1, 2.  
 1893 *Phylloporina angulata* (Hall); Foerste, p. 600, pl. 28, fig. 1.  
 1919 *Chasmatopora angulata* (Hall); Foerste, p. 369.  
 2010a *Chasmatopora foerstei* McKinney and Wyse Jackson, p. 444, text-fig. 1A–F.  
 2010b *Chasmatopora foerstei* McKinney and Wyse Jackson, p. 41.

*Holotype.*—USNM 84851 (Foerste, 1893, fig. 1). Clinton Group, Silurian; Soldiers' Home Quarries, Dayton, Ohio, USA.

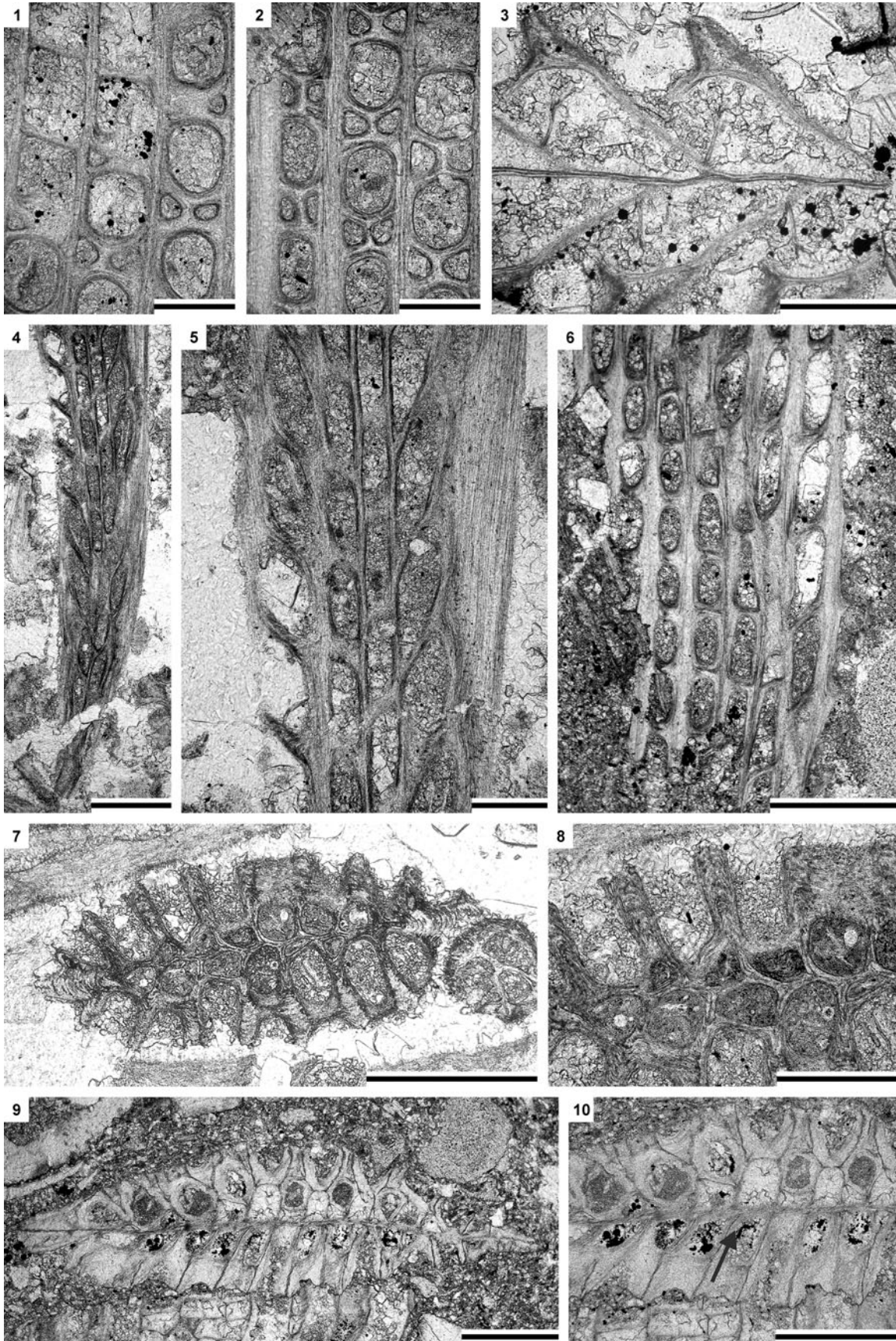
*Occurrence.*—Silurian (Llandovery); Ohio and New York, USA.

*Description.*—Reticulate colony consisting of anastomosing branches with regular oval to lens-shaped fenestrules. Branches rounded to triangular in transversal section, 0.25–0.45 mm wide. Autozoecia long, having oblong-rectangular shape in deep tangential section, well-developed vestibules, arranged in three to four alternating rows on branches. Hemisepta absent. Diaphragms common, usually situated near the bend of autozoecia in exozone. Terminal diaphragms not observed. Autozoecial apertures oval, having distinct peristomes, surrounded by 8–10 nodes. Apertural nodes 0.010–0.015 mm in diameter. Keels absent. Nodes moderately large, regularly spaced on the obverse side. Internal granular skeleton well developed, 0.015–0.020 mm thick, continuous with medial laminae, peristomes, and microstyles. Outer laminated skeleton well developed, 0.08–0.09 mm thick. Reverse side striated, containing abundant microstyles. Microstyles 0.015–0.030 mm in diameter. Heterozoecia not observed.

*Materials.*—SMF 23.710–SMF 23.733.

*Remarks.*—*Chasmatopora foerstei* McKinney and Wyse Jackson, 2010a differs from *C. silurica* (Kopajevich, 1975) from the Silurian (Llandovery) of Ukraine in its narrower branches (branch width 0.25–0.40 mm versus 0.59–0.63 mm in *C. silurica*). *Chasmatopora foerstei* is similar to *C.*

**Figure 11.** (1–10) *Moyerella parva* n. sp.: (1–3) longitudinal section of a colony segment with a pointed base and widened proximal part showing medial axis and autozoecia, holotype SMF 23.559; (4) tangential section showing autozoecial apertures, tubules, and tectitooecia, holotype SMF 23.559; (5) longitudinal section of a colony showing medial axis and autozoecia, paratype SMF 23.580; (6) longitudinal section of a colony showing shape of autozoecia, paratype SMF 23.575; (7) tangential section showing autozoecial apertures, tubules, and metazoecia (arrows), paratype SMF 23.578; (8) tangential section showing autozoecial apertures, tubules, and acanthostyles (arrows), paratype SMF 23.621; (9, 10) tangential section showing autozoecial apertures and acanthostyles, paratype SMF 23.602. (11–14) *Phaenopora multifida* (Hall, 1883): (11, 12) tangential section showing autozoecial apertures and metazoecia divided by longitudinal crests, SMF 23.671; (13, 14) branch transverse section showing mesotheca, autozoecia, and metazoecia, SMF 23.670. (1–3, 5, 6, 9, 12, 13) Scale bars = 0.5 mm; (4, 7, 8, 10, 14) scale bars = 0.2 mm; (11) scale bar = 1 mm.



**Table 8.** Descriptive statistics of *Phaenopora multifida* (Hall, 1883). N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Branch width	13	2.07	0.803	38.81	1.00	3.50
Branch thickness	13	0.60	0.201	33.35	0.35	1.15
Aperture width	20	0.08	0.006	7.12	0.08	0.10
Aperture spacing along branch	20	0.26	0.024	9.23	0.19	0.31
Aperture spacing diagonally	20	0.21	0.020	9.81	0.18	0.25
Metazooecia width	20	0.030	0.005	17.52	0.025	0.040

*granistriata* (Ulrich, 1890) from the Ordovician of the United States and Canada but differs from it in narrower branches (branch width 0.25–0.40 mm versus 0.30–0.50 mm in *C. granistriata*). Moreover, *C. granistriata* possesses keels on the obverse colony surface, which are absent in the present species.

Family Fenestellidae King, 1849  
Genus *Fenestella* Lonsdale, 1839

*Type species.*—*Fenestella subantiqua* d'Orbigny, 1850. Lower Silurian (Wenlockian); England.

*Fenestella jupiterensis* Bassler, 1928  
Figure 14.4–14.7; Table 13

1928 *Fenestella jupiterensis* Bassler, p. 157, pl. 11, fig. 14, pl. 13, figs. 2, 3.

*Holotype.*—USNM PAL 69057. Jupiter Formation, Llandoverly, Silurian; Anticosti, Canada.

*Occurrence.*—Jupiter Formation, Llandoverly, lower Silurian; Canada. Reynales Formation, Hickory Corners Member, Aeronian, Llandoverly, lower Silurian; Hickory Corners, New York, USA.

*Exterior description.*—Reticulate colony formed by straight branches joined by wide dissepiments. Fenestrules oval to rectangular. Autozoecia arranged in two rows on branches. Autozoecial apertures circular, with low peristome; 10–12 nodes in peristome, 0.008–0.010 mm in diameter; three to four apertures spaced per fenestrule length. Keel low, without nodes. Reverse side bearing nodes.

*Interior description.*—Autozoecia relatively long, roughly rectangular in mid-tangential section with moderately long vestibule in longitudinal section. Axial wall between autozoecial rows straight; aperture positioned at distal end of chamber. Hemisepta absent. External laminated skeleton well developed on both obverse and reverse sides traversed by

**Table 9.** Descriptive statistics of *Ptilodictya sulcata* Billings, 1866. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Aperture width	20	0.07	0.005	6.44	0.07	0.08
Aperture spacing along branch	20	0.25	0.025	9.76	0.22	0.31

microstyles. Microstyles 0.010–0.015 mm in diameter. Heterozooecia not observed.

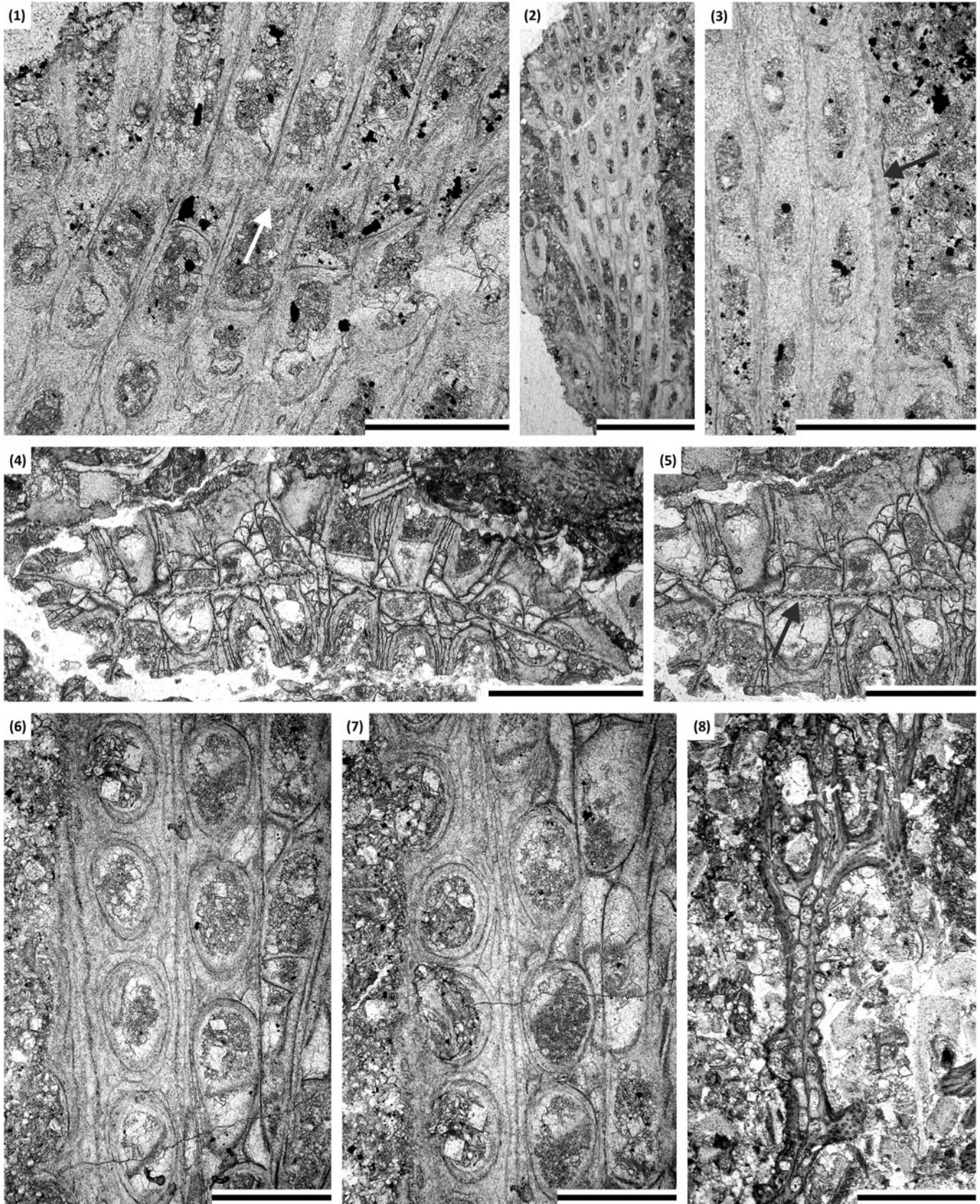
*Materials.*—SMF 23.734–SMF 23.755.

*Remarks.*—*Fenestella jupiterensis* Bassler, 1928 differs from *F. elegans* Hall, 1852 from the Rochester Shale (New York) in absence of nodes on the keel and spacing of three to four apertures per fenestrule instead four to five as in the latter species. *Fenestella jupiterensis* is similar to another species from the Jupiter Formation, *F. anticostiensis* Bassler, 1928. Unfortunately, all these species are described very briefly, without thin sections and detailed measurements. *F. anticostiensis* also possesses medial keel without nodes but differs in slightly larger dimensions. Bassler (1928, p. 156–157) gave only the numerical parameter for the spacing of fenestrules: “3 fenestrules in 3 mm longitudinally and 4 in the same space transversely.” These measurements produce estimates for average values of the distance between dissepiment centers (1 mm versus 0.73 mm in the present species) and of the distance between branch centers (0.75 mm versus 0.46 mm in the present species).

## Discussion

*Ecological interpretation of the fauna.*—The bryozoan fauna from the Reynales Formation (lower Silurian, Aeronian) at Hickory Corners, near Lockport, New York, contains 13 species: one cyclostome (*Diploclema argutum* Bassler, 1906), two cystoporates (*Cheilotrypa* aff. *C. variolata* [Hall, 1876] and *Hennigopora apta* Perry and Hattin, 1960), three trepostomes (*Homotrypa niagarensis* n. sp., *Hallopora* aff. *H. elegantula* [Hall, 1852], and *Leioclema adsuetum* n. sp.), five cryptostomes (*Moyerella parva* n. sp., *Phaenopora multifida* [Hall, 1883], *Ptilodictya sulcata* Billings, 1866, *Metadictya* sp., and *Trigonodictya* sp.), as well as two fenestrates (*Chasmatopora foerstei* McKinney and Wyse Jackson, 2010a and *Fenestella jupiterensis* Bassler, 1928). This fauna is distinguished by the dominance of erect growth forms of different sizes and configurations. *Hennigopora apta* is the only strictly encrusting species, whereas *Leioclema adsuetum* is observed to have produced both encrusting and erect branched colonies, often with secondary overgrowths (Fig. 10.2, 10.3). *Hallopora* aff. *H. elegantula* (Hall, 1852) is

**Figure 12.** (1–3) *Phaenopora multifida* (Hall, 1883): (1, 2) tangential section showing autozoecial apertures and metazooecia, SMF 23.671; (3) longitudinal section showing superior and inferior hemisepta, SMF 23.638. (4–8) *Ptilodictya sulcata* Billings, 1866: (4, 5) branch oblique section showing autozoecial chambers, SMF 23.693; (6) tangential section showing autozoecial apertures, SMF 23.694; (7, 8) branch transverse section showing autozoecial chambers and mesotheca, SMF 23.691. (9, 10) *Metadictya* sp.: branch oblique section showing autozoecial chambers and mesotheca, SMF 23.703. (1–3, 5, 8) Scale bars = 0.2 mm; (4, 6, 7, 9, 10) scale bars = 0.5 mm.



**Table 10.** Descriptive statistics of *Metadictya* sp. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Aperture width	10	0.07	0.008	11.66	0.06	0.08
Aperture spacing along branch	10	0.37	0.021	5.78	0.34	0.40
Aperture spacing diagonally	10	0.24	0.017	7.30	0.20	0.26

an erect branched species that also developed secondary overgrowths (Fig. 9.2, 9.3). *Homotrypa niagarensis* produced robust branched colonies with extremely rare secondary overgrowths.

*Cheilotrypa* aff. *C. variolata* produced erect hollow, pseudotubular colonies. Separate encrusting sheets of this bryozoan are extremely rare, apparently present only in its basal parts. However, it can be assumed that this species is not truly tubular but represents an encrustation of a soft-bodied, erect branching organism (likely algae) by bioimmuration. After decay of the encrusted organism, the rigid bryozoan colony retained its shape. The interiors of wider tubes are usually filled by micrite (Figs. 6.4, 6.5, 6.7, 6.8, 7.1, 7.5), whereas narrower tubes are commonly filled by sparry calcite (Fig. 6.6). Presence of micrite implies access to the seawater with sediment particles, which means that the larger tubes were not completely encrusted by bryozoans. By contrast, smaller individuals of the soft-bodied organism were probably completely encrusted by bryozoans. Locally, contact between the bases of the encrusting sheets and the substrate is observed (Figs. 6.7, 6.8, 7.5). Longitudinal sections reveal regularly narrowing portions, which apparently display the form of the encrusted organism. This encrustation may have been obligatory because no other objects (except a few sheets on the substrate) are encrusted by this bryozoan.

The nature of the bioimmured organism remains unclear. Regular narrowings are typical of algae and sponges. In both cases, the bioimmuration must have been detrimental as access to seawater and light would have been interrupted. The majority of tube lumens are rounded or oval, but many are also narrowly ovoid (Figs. 6.8, 7.1). That may represent the original shape of the soft-bodied organism or could have occurred due to collapse of the bryozoan tube after the decomposition of the encrusted host.

Hollow tubular or hollow cylindrical growth forms were called ‘pseudovinculariform’ by Stach (1936) and are typically the result of encrusting forms that completely surround the stem of a host plant or animal substrate (Hageman et al., 1998). Mature colonies are able to continue their growth as erect arborescent forms with hollow branches (Hageman et al., 1998). Bryozoans encrust various soft-bodied organisms of cylindrical shape, such as algae, sponges, ascidians, and octocorals. Encrustation of ephemeral substrates is very common in modern cool-water carbonates but apparently less common in pre-Cretaceous bryozoan-rich environments (Hageman et al.,

**Table 11.** Descriptive statistics of *Trigonodictya* sp. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Aperture width	10	0.20	0.019	9.30	0.18	0.24
Aperture spacing along branch	9	0.46	0.031	6.65	0.42	0.50

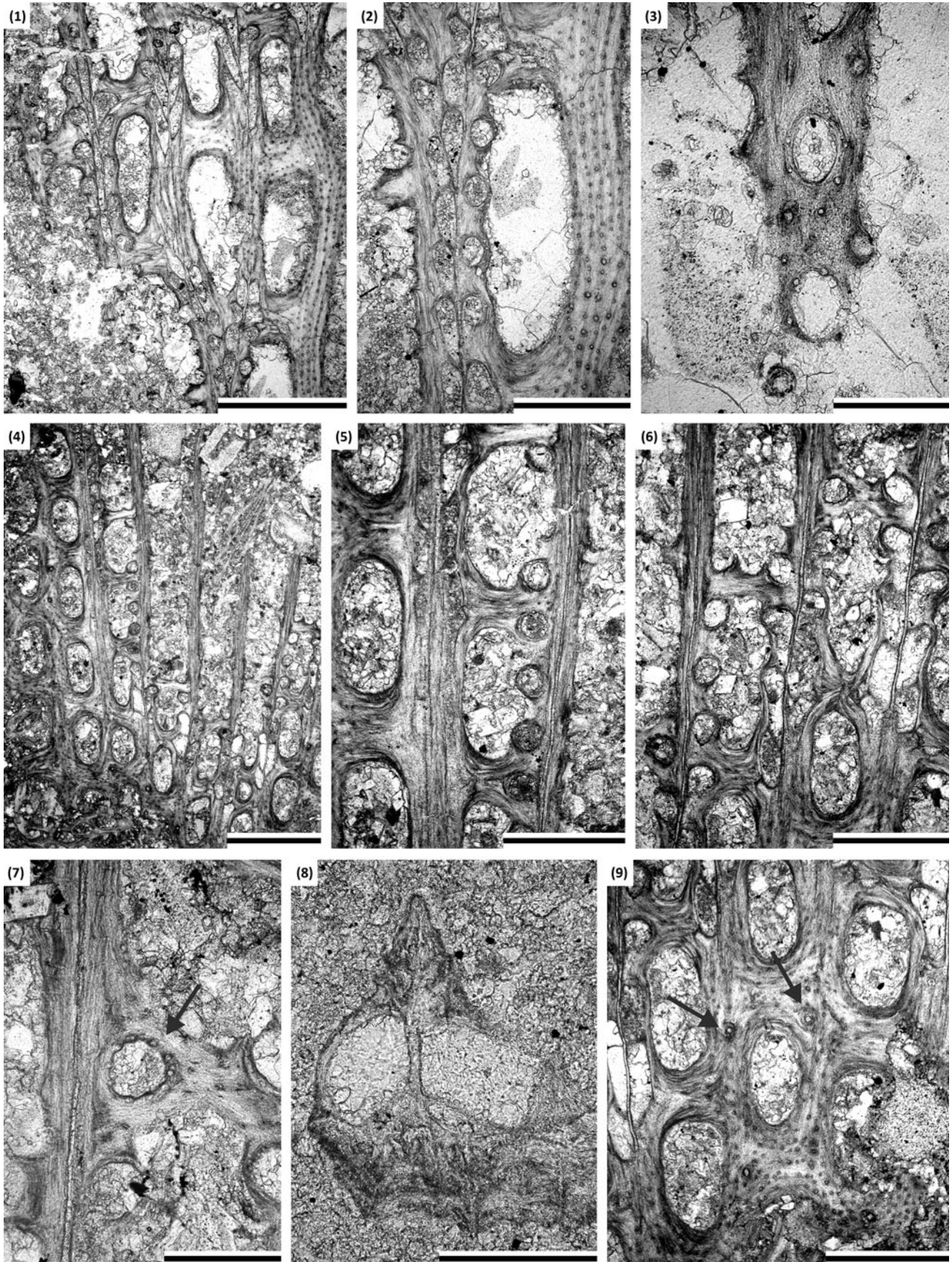
2000). However, hollow cylindrical colonies repeatedly occurred among Paleozoic bryozoans, notably in cystoporates and trepostomes, which potentially develop encrusting colonies. Representatives of the orders Fenestrata and Cryptostomata developed mainly arborescent colonies of various configurations. However, exceptions are known in both of these orders. For example, cryptostome genus *Rhabdomeson* developed obligatory cylindrical colonies, with a central cylinder that is considered to be a polymorph (Blake, 1976, 1983). In fenestrates, the genus *Bigeyina* is known to form hollow conical colonies (Suárez-Andrés and McKinney, 2010).

Hollow cylindrical colonies of *Cheilotrypa ostiolata* (Hall, 1852) are abundant in the younger Rochester Shale (Wenlock) of New York (Bassler, 1906). Longitudinal sections of this bryozoan show regular narrowings similar to those in material from the Reynales Formation (Bassler 1906, pl. 9, fig. 4). However, the voids in the Rochester Shale bryozoan are narrower than those in *Cheilotrypa* aff. *C. variolata*: 0.22–0.35 mm versus 0.42–1.35 mm. *Cheilotrypa ostiolata* (Hall, 1852) from the Rochester Shale also appears to have been an obligatory encruster of its host substrates, with minor development of freely encrusting sheets. Arborescent colonies have advantages over encrusting bryozoans in their higher tiering (and consequently, better feeding and reproductive capacity) and escape from competition for substrate (McKinney and Jackson, 1989). Otherwise, it has been observed that sessile animals tend to avoid erect growth in the presence of predators and switch to encrusting mode of growth (e.g., Guida, 1976; McKinney and Jackson, 1989). Absence of predation in the Silurian, in addition to other ecological factors, may also have favored development of hollow cylindrical colonies instead of simply encrusting ones. The flexibility of bryozoans in developing various growth forms depending on their ecological situation (e.g., Harmelin, 1976) is also a strongly confusing factor for taxonomists who can misinterpret growth form as taxonomic character, which could be utilized for discrimination of species or even genera.

Cryptostome bryozoans are represented by one rhabdomesid species and four ptilodictyines. *Moyerella parva* developed an articulated colony consisting of flexibly connected segments. Separate segments have tapered bases and widened apical parts representing ‘sockets.’ Such colonies are ubiquitous among arthrostyloid cryptostomes (e.g., Wyse Jackson et al., 2017) and even more common in Cenozoic cyclostomes and cheilostomes (Taylor and James, 2013). Stach (1936) named such colonies ‘cellariiform,’ after the cheilostome bryozoan *Cellaria*. This

**Figure 13.** (1–3) *Metadictya* sp.: (1) branch oblique section showing autozooeical chambers and mesotheca with tubules (arrow), SMF 23.695; (2, 3) branch oblique section showing autozooeical apertures and spherules (arrow), SMF 23.695. (4–7) *Trigonodictya* sp.: (4, 5) branch transverse section showing autozooeical chambers, vesicles, and mesotheca with tubules (arrow), SMF 23.706; (6, 7) tangential section showing autozooeical apertures and vesicles, SMF 23.708. (8) *Chasmatopora foerstei* McKinney and Wyse Jackson, 2010a: tangential section showing branches with autozooeica, SMF 23.730. (1) Scale bar = 0.2 mm; (2, 4, 8) scale bars = 1 mm; (3, 5–7) scale bars = 0.5 mm.





**Table 12.** Descriptive statistics of *Chasmatopora foerstei* McKinney and Wyse Jackson, 2010a. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Branch width	20	0.35	0.042	12.03	0.25	0.40
Dissepiment width	16	0.28	0.042	14.91	0.19	0.34
Aperture width	20	0.07	0.009	12.68	0.06	0.09
Aperture spacing along branch	20	0.25	0.015	5.96	0.22	0.27
Aperture spacing diagonally	20	0.18	0.019	10.35	0.14	0.21
Node width	15	0.046	0.006	12.97	0.038	0.055

type of the colony form was initially assumed to be adapted for the life in agitated, rather shallow waters; however, such forms have also been found in depths up to 250 m in the modern (e.g., Bone and James, 1993; Moissette, 2000; Amini et al., 2004).

Four ptilodictyine species developed bilaminar (bifoliate) colonies in which zooecia open on both sides of a flat, lenticular branch (e.g., Figs. 11.13, 12.7, 12.9, 13.4). Ptilodictyines were an extremely successful group in the Ordovician and to a similar extent in the Silurian, but they declined in the middle Paleozoic and disappeared in the early Permian (Karklins, 1983; Gorjunova and Lavrentjeva, 1993; Ernst and Nakrem, 2007). *Ptilodictya sulcata* from the described fauna developed a lanceolate colony with a tapering proximal segment (Fig. 12.4) that was apparently flexibly joined to a socket-shaped base, which is usual in *Ptilodictya* (Hennig, 1905; Ross, 1960a, b). Three other ptilodictyine species seem to have been firmly attached to the substrate.

The majority of studies devoted to the distribution of bryozoan growth forms with depth have been conducted on modern faunas. These interpretations may not be precisely analogous to similar Paleozoic growth forms, but the general trends may be similar (Taylor, 2005). The modern bryozoan faunas, for example, show an increasing ratio of erect species with depth (e.g., Schopf, 1969; Schopf et al., 1980; McKinney and Jackson, 1989; Moissette, 2000; Amini et al., 2004).

The common usage of the term ‘fenestrate’ appears confusing in regard to reticulate, net-like colonies produced by branches jointed by dissepiments or fused together (anastomosing). In the taxonomic sense, ‘fenestrate’ means belonging to the order Fenestrata, which includes a variety of forms and is restricted to the Paleozoic. For the description of such colonies, the terms ‘reticulate’ or ‘net-like’ are more appropriate and concern rather the form than the taxonomic assignment. Reticulate colonies appeared convergently in various bryozoan groups in both stenolaemate and gymnolaemate clades, presumably because they are efficient for filtering (e.g., Cowen and Rider, 1972; McKinney, 1977; McKinney and Jackson, 1989; Taylor and James, 2013). Bryozoans with reticulate colony shapes are regarded as being adapted to various types of environments, although generally avoiding shallow high-energy biotopes

**Table 13.** Descriptive statistics of *Fenestella jupiterensis* Bassler, 1928. N = number of measurements; X = mean; SD = sample standard deviation; CV = coefficient of variation; Min = minimum value; Max = maximum value; units = mm.

	N	X	SD	CV	Min	Max
Branch width	25	0.27	0.031	11.4	0.22	0.35
Dissepiment width	30	0.22	0.039	18.2	0.16	0.31
Fenestrate width	23	0.21	0.035	17.1	0.16	0.30
Fenestrate length	30	0.50	0.063	12.6	0.32	0.65
Distance between branch centers	30	0.46	0.066	14.4	0.35	0.59
Distance between dissepiment centres	30	0.73	0.079	10.8	0.48	0.84
Aperture width	39	0.09	0.006	6.9	0.08	0.10
Aperture spacing along branch	40	0.22	0.023	10.2	0.18	0.26
Aperture spacing diagonally	10	0.22	0.023	10.8	0.20	0.28
Maximal chamber width	17	0.10	0.008	7.9	0.09	0.11
Node width	17	0.053	0.007	12.6	0.040	0.065

(e.g., McKinney and Gault, 1980; Kelly and Horowitz, 1987; Nelson et al., 1988; Amini et al., 2004).

Bryozoans of the Reynales Formation show medium robustness of their colonies. They can be divided into delicate (< 2 mm in thickness/diameter) and robust (> 2 mm) forms. The former category includes six species: *Diploclema argutum*, *Hennigopora apta*, *Moyerella parva*, *Ptilodictya sulcata*, *Chasmatopora foerstei*, and *Fenestella jupiterensis*. The latter category includes seven species: *Cheilotrypa* aff. *C. ariolate*, *Homotrypa niagarensis*, *Hallopora* aff. *H. elegantula*, *Leioclema adsuetum*, *Phaenopora multifida*, *Metadictya* sp., and *Trigonodictya* sp. The robustness can be considered as a factor of depth. In general, deeper faunas contain species with more delicate colonies (e.g., Thomsen, 1977; Schopf et al., 1980; Amini et al., 2004).

Summarizing the colony shapes and their robustness, we conclude that the fauna from the Reynales Formation at Hickory Corners existed on stable substrate under moderate wave energy conditions. Scarcity of encrusting forms as far as relatively high proportion of delicate colonies indicates a relatively deeper and quiet environment. The low level of fragmentation of organic remnants (Figs. 4.1, 8.1) supports the absence of strong currents. However, the presence of thick-branched trepostomes and such lithological characteristics as well-washed sparry grainstones (Fig. 4.2) points to relatively high wave activity. Flexibly jointed colonies (*Moyerella*, *Ptilodictya*) are adapted for both kinds of biotopes. The presence of large branched trepostomes and lenticular cryptostomes suggests significant stability of the substrate. In general, the bryozoan fauna correlates well with the interpretation made by Brett et al. (1993) for this area as belonging to Benthic Assemblage 4 situated in the shallow (30–60 m depth) outer shelf environment.

*Species richness within samples.*—Paleozoic bryozoans have an immense advantage over many other fossils in that they can be identified even as small fragments because of their modularity and specific skeletal material (mainly low-magnesium calcite). Their study must be undertaken with use of thin sections. In

**Figure 14.** (1–3) *Chasmatopora foerstei* McKinney and Wyse Jackson, 2010a: tangential section showing autozooecial chambers and apertures with nodes, nodes between autozooecial apertures, and microstyles on the reverse side of branches, SMF 23.710. (4–9) *Fenestella jupiterensis* Bassler, 1928: (4–7) tangential section showing autozooecial chambers and apertures with nodes (arrow) and keel without nodes, SMF 23.740; (8) branch transverse section, SMF 23.745; (9) tangential section showing reverse side of branch with nodes (arrows) and microstyles, SMF 23.740. (1, 4) Scale bars = 1 mm; (2, 5, 6, 9) scale bars = 0.5 mm; (3, 7, 8) scale bars = 0.2 mm.

case the isolation of separate colonies from solid rock is impossible, preparation of serial thin sections appears the best method for study. Collected material from the Reynales Formation included ~40 fist-sized (~10 × 10 cm) rock samples from which 20 samples were used for preparation of thin sections (11 thin sections of 24 × 48 mm size and 67 thin sections of 50 × 50 mm size). All samples were taken from the same stratigraphical and ecological level alongside the outcrop. Thin sections were oriented both in plane of bedding and perpendicular to it. The number of thin sections varied from 1 to 11 per sample. On average, 6 bryozoan species occurred per large thin section (50 × 50 mm object glass, roughly 16 square cm of rock surface), varying from 3 to 8. The number of species per sample varied from 6 to 11, averaging 8 species per sample (Table 14). The richness of bryozoans generally increased with the number of thin sections per sample, but not in all cases. Sample HC-23 includes 3 thin sections that contain a total of 11 species, whereas the sample HC-19 contains 8 species in 8 thin sections.

*Diploclima argutum*, *Cheilotrypa* aff. *C. variolata*, *Moyerella parva*, and *Phaenopora multifida* occur in almost every thin section. *Chasmatopora foerstei* and *Fenestella jupiterensis* are also very common, occurring in more than 90% of thin sections. *Hallopora* aff. *H. elegantula* occurs in 70% of samples, and *Leioclema adsuetum* is present in 50% of samples. *Hennigopora apta*, *Ptilodictya sulcata*, *Metadictya* sp., and *Trigonodictya* sp. are uncommon, occurring in less than 25% of samples.

*Moyerella parva* is the most abundant species by the number of fragments found in the thin sections. This can be explained as a taphonomic effect. *Moyerella parva* possessed an articulated colony, which apparently consisted of several segments connected by flexible organic joints. After death and decomposition of organic material of joints, the segments were disseminated in the sediment. Therefore, the number of segments should be higher than the number of individual colonies.

**Biogeography.**—The studied bryozoan fauna contains mainly cosmopolitan genera such as *Hallopora*, *Homotrypa*, *Leioclema*, and *Phaenopora*. Outside North America, *Diploclima* is known in Europe (Gotland, Great Britain, and Estonia). This genus might have a wider distribution, however, as it can be easily overlooked because of its small size. Silurian species of *Cheilotrypa* are mainly restricted to North America. By contrast, *Moyerella* was previously known only from Estonia and Russia. *Hennigopora* is temporally restricted to the Silurian; geographically, however, the lower Silurian species of *Hennigopora* are known from North America, China, and Russia. *Metadictya* is known from the Silurian of North America and Europe (Sweden and Estonia). Silurian species of *Trigonodictya* are restricted to North America, whereas the Silurian *Ptilodictya* species are known from North American and European localities as well as from Siberia.

The identified species have rather local relations except *Diploclima argutum* (Bassler, 1906), which was also identified in Europe (Sweden and Estonia). *Hennigopora apta* Perry and Hattin, 1960 was originally described from the Osgood Formation (Telychian-Sheinwoodian) of Indiana, USA. *Phaenopora multifida* (Hall, 1883) is known from the Clinton Group of Ohio. *Chasmatopora foerstei* McKinney and Wyse Jackson,

**Table 14.** Species distribution in the studied material (78 thin sections of 20 rock samples). 1 = *Diploclima argutum*; 2 = *Cheilotrypa* aff. *C. variolata*; 3 = *Hennigopora apta*; 4 = *Homotrypa niagarensis*; 5 = *Hallopora* aff. *H. elegantula*; 6 = *Leioclema adsuetum*; 7 = *Moyerella parva*; 8 = *Phaenopora multifida*; 9 = *Ptilodictya sulcata*; 10 = *Metadictya* sp.; 11 = *Trigonodictya* sp.; 12 = *Chasmatopora foerstei*; 13 = *Fenestella jupiterensis*.

	1	2	3	4	5	6	7	8	9	10	11	12	13
HC-1	x	x	—	x	—	—	x	x	—	—	—	x	x
HC-6	x	x	—	—	—	x	x	x	—	—	—	x	—
HC-7	—	x	—	x	—	x	x	x	—	—	x	x	—
HC-8	x	x	—	—	x	—	x	x	—	—	x	x	x
HC-9	x	x	—	—	—	—	x	x	—	x	—	x	x
HC-10	x	x	—	—	x	—	x	x	—	—	—	x	x
HC-11	x	x	x	—	x	—	x	x	—	x	—	x	x
HC-13	x	x	—	x	x	x	x	x	x	—	—	x	x
HC-16	x	x	—	—	—	x	x	x	—	—	—	x	x
HC-17	x	x	—	—	—	—	x	x	—	—	—	x	x
HC-18	x	x	—	x	x	—	x	x	—	x	—	x	x
HC-19	x	x	—	—	x	—	x	x	x	—	—	x	x
HC-20	x	x	x	—	x	x	x	x	—	—	x	x	x
HC-21	x	x	x	x	x	x	x	x	—	—	—	x	x
HC-22	x	x	—	x	x	x	x	x	—	x	x	x	—
HC-23	x	x	—	x	x	x	x	x	x	—	x	x	x
HC-25	x	x	—	x	x	x	x	x	—	—	—	x	x
HC-26	x	x	—	x	x	—	—	x	—	—	—	x	x
HC-27	x	x	x	—	x	x	x	x	—	—	—	—	x
HC-28	x	x	—	x	x	—	x	—	—	x	—	—	x

2010a is known from different localities of the Clinton Group in Ohio and New York. *Fenestella jupiterensis* Bassler, 1928 was originally described from the Jupiter Formation (Llandoverly, Telychian, lower Silurian) of Anticosti, Canada. *Ptilodictya sulcata* Billings, 1866 is known from the Upper Ordovician to lower Silurian succession of Anticosti. Furthermore, this species is present in the Upper Ordovician of Estonia (Bassler, 1911) and in erratic boulders of Poland (Kiepur, 1962).

## Conclusions

Bryozoan fauna from the Brewer Dock (Hickory Corners) Member of the Reynales Formation (lower Silurian, Aeronian) at the type locality, Hickory Corners on Lockport Junction Road, in western New York, USA, is abundant and moderately diverse. The studied assemblage contains 13 species, from which 3 are new: trepostomes *Homotrypa niagarensis* n. sp. and *Leioclema adsuetum* n. sp. and the rhabdomesine cryptostome *Moyerella parva* n. sp. This fauna is clearly dominated by erect growth forms of various shapes and robustness. The minority of encrusting forms (one obligatory species) and equal ratio of delicate and robust forms suggest a moderately agitated environment with a stable substrate. Bryozoans are abundant and diverse within separate samples, with an average of six species occurring within a single thin section of ~16 cm<sup>2</sup> surface area. The species composition reveals paleobiogeographic connections to other Silurian localities of New York as well as Ohio and Indiana (USA) and Anticosti (Canada), whereas the generic composition suggests connections with bryozoan faunas of Laurentia and Baltica.

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