

23 Rhaetian, thus constraining its deposition to a time interval characterized by increasing global
24 humidity and seasonality. The integrated palynofacies and lithofacies data enabled characteri- zation
25 of the timing of the drowning phases of the carbonate platform-basin system as being controlled by
26 rela- tive sea level changes mostly triggered by the Triassic extensional tectonic activity. During the
27 first phase of the relative sea-level rise, clayey and organic-rich sediments were deposited only in the
28 deepest portion of the basin. As the sea level continued to rise, the entire system drowned completely
29 and suboxic-anoxic basinal sediments were deposited across the whole Hyblean region, onlapping the
30 shallow-water facies. In the meantime increasing global humidity contributed to an increased
31 freshwater input in the marine depositional system as documented by the presence of fern spores and
32 clay. It caused water stratification and subsequent anoxia at marine basins, fa- voring the preservation
33 of sedimentary organic matter. This atmospheric change could be related to the degassing of the
34 Central Atlantic Magmatic Province.

35 *Key words: palynofacies; palynostratigraphy; platform-basin system; Late Triassic; Sicily (Italy)*

36

37 **1. Introduction**

38 The temporal and spatial distribution of Phanerozoic organic rich sediments relates to a combination
39 of variables: organic productivity, appropriate sedimentation rates and organic matter preservation
40 (Tyson, 2001, 2005; Katz, 2005; Trabucho-Alexandre et al., 2012). Organic carbon rich sediments
41 have been commonly attributed to widespread ocean anoxia and to associated water column
42 stratification that isolates the marine bottoms from the oxygen mixed zone (Tyson and Pearson, 1991;
43 Wignall and Newton, 2001; Pancost et al., 2004; Harris, 2005; Meyer and Kump, 2008). Other
44 theories suggest that a combination of high primary productivity and low sedimentation rate promotes
45 the accumulation of large amounts of organic matter due to a relatively low degree of dilution by
46 siliciclastics and skeletal debris (Lallier-Vergès et al., 1995; Perkins et al., 2008). During the Late

47 Triassic – Early Jurassic times in the western Tethys realm, deposition of organic rich clay and marly
48 successions seems to coincide with a climate warming and an increasing rainfall and runoff (Korte et
49 al., 2009; Bonis et al., 2010; Haas et al., 2010, 2012; Michalík et al., 2010; Berra, 2012). This has
50 been interpreted as a consequence of an intense monsoonal activity (Parrish, 1993; Satterley, 1996;
51 Sellwood and Valdes, 2007; Bonis and Kürschner, 2012). In addition, the onset of igneous and
52 volcanic activity within the Central Atlantic Magmatic Province (CAMP), is generally believed to
53 have strongly influenced the climate change by releasing of volcanic gases (mainly CO₂ and SO₂) into
54 the ocean–atmosphere system (Marzoli et al. 2004; Cirilli et al., 2009; van de Schootbrugge et al.,
55 2009; Lucas et al., 2011; Ruhl et al., 2011; Schaller et al., 2011; Pálffy and Zajzon, 2012; Vajda et al.,
56 2013; Bond and Wignall, 2014; Lindström, 2016; Davies et al., 2017; Lindström et al., 2017b).
57 Consequently, the end-Triassic mass extinction (ETE) has been attributed to the huge amount of
58 greenhouse-gas emissions in the atmosphere and in the ocean waters and associated to the ≈3-6%
59 negative carbon isotope excursion, recorded in both terrestrial and marine environments (Schoene et
60 al., 2010; Whiteside et al., 2010; Ruhl et al., 2011; Dal Corso et al., 2014; Lindström, 2016;
61 Lindström et al., 2017b). Recent high-precision U-Pb ages from CAMP mafic intrusive units (Davies
62 et al., 2017) and large scale correlations based on a set of integrated biotic, geochemical and
63 radiometric data (Lindström et al., 2017b), document that magmatic activity started about 100 Kyr
64 before the earlier known eruptions, providing evidence of the causal relation between CAMP and
65 ETE. The resulting increase of fresh water supply and nutrient input in the marine environments, via
66 river runoff, led to water density stratification and increased primary productivity that drove the
67 organic matter accumulation and preservation at the anoxic marine bottoms.

68 The overall aim of this study is to provide new insights into the factors triggering the accumulation
69 and preservation of organic matter within a given paleoclimatic and paleogeographic scenario related
70 to the well-known source and seal rocks of the Hyblean Petroleum System, in south eastern Sicily

71 (Italy). We combined palynofacies and lithofacies analyses of two on-shore wells from Sicily in order
72 to interpret the paleoenvironmental and paleoclimatic conditions and the input of organic matter
73 preserved in the sediments. The data provided in this paper complement and integrate those presented
74 by Cirilli et al. (2015) regarding another on-shore well (Pachino 4) drilled in the same area.

75

76 **2. Geological setting**

77 In the last few decades, Eni Upstream and Technical Services have carried out many geological
78 studies, in southeastern Sicily on-shore and off-shore, following exploration activity (Fig. 1) (Frixia et
79 al., 2000; Trincianti et al., 2015 for references). The studied wells are located in the south east of
80 Sicily (Hyblean Plateau) which is characterized by an over 5 km thick Triassic-Neogene sequence,
81 lying above a 20–25 km thick sequence with African affinity and acting as a foreland basin during
82 Neogene Alpine orogenesis (Patacca et al., 1979; Yellin-Dror et al., 1997; Catalano et al., 2000,
83 2002, 2013; Granath and Casero, 2004; Finetti et al., 2005). Small and large-scale paleogeographic
84 reconstructions reveal that Sicily has been located along the African - European plate boundary from
85 Paleozoic times (Ruiz-Martínez et al., 2012; Catalano et al., 2013; Berra and Angiolini, 2014; Scotese
86 and Schettino, 2017). Starting from Triassic, the west-central Mediterranean platform (Apulia-Adria)
87 was broken apart by the same tectonic processes that caused the opening of the Central Atlantic.
88 After separation of north west Africa (northern Gondwana) from Eastern North America, Adria,
89 Apulia (including Sicily) and southern Turkey continued to be part of the African Plate (Catalano et
90 al., 2002, 2013; Robertson et al., 2003; Finetti et al., 2005; Berra and Angiolini, 2014; Scotese and
91 Schettino, 2017). The extensional faulting was accompanied by regional and large-scale fissural
92 basaltic volcanism at least since the Triassic. Intercalations of mafic volcanics have been recorded in
93 several wells of the Hyblean Plateau at different stratigraphic levels (Patacca et al., 1979; Rocchi et
94 al., 1996; Granath and Casero, 2004; Finetti et al., 2005; Scotese and Schettino, 2017). Beginning in

95 the Late Triassic, the whole Hyblean Plateau region was occupied by a wide shallow water carbonate
96 platform (represented by the Sciacca Formation). The extensional phase related to the continental
97 rifting caused the carbonate platform to break-up and this triggered the onset of a platform to basin
98 system (Patacca et al., 1979; Brosse et al., 1988; Frixia et al., 2000). Shallow-water carbonate deposits
99 (Noto Formation) covered the northern part of the area, while a deep anoxic-suboxic intraplatform
100 basin developed southward (Fig. 1). The Noto Formation (about 300 m thick) includes at least three
101 interfingering facies. The first facies is spread throughout the Hyblean Plateau and consists of
102 limestones (mudstones and wackestones), often dolomitized and recrystallized, interlayered with
103 organic rich black shales. The second facies, is only found at the edge of the Plateau, is composed of
104 wackestones, packstones and oolitic grainstones interpreted as beach ridge deposits (Patacca et al.,
105 1979; Brosse et al., 1988). have been considered as beach ridge deposits (Patacca et al., 1979; Brosse
106 et al., 1988). The third facies, named the Mila Member of the Noto Formation, occurs in the marginal
107 area of the carbonate platform (Fig. 2). The Mila Member consists of two superimposed carbonate
108 bodies, which backstep northwards, and are locally recrystallized and dolomitized. The lower
109 microbial body lies above the carbonate platform of the Sciacca Formation, while the upper body
110 overlies the lower microbial body (basinwards) and the Noto Formation (landwards, Frixia et al.,
111 2000; Felici et al., 2014) (Fig. 2). As subsidence increased, sedimentation in the inner platform was
112 mostly dominated by limestones and organic rich shales (still included in the Noto Formation; Frixia
113 et al., 2000). At the same time, in the rapidly subsiding adjacent basin, a thick succession
114 (Streppenosa Formation) deposited under suboxic-anoxic conditions. The thickness of the
115 Streppenosa Formation is variable and reaches a maximum of about 3000 m in the southeastern part
116 of the Hyblean Plateau. It has been subdivided into three members (Frixia et al., 2000). The Lower
117 Streppenosa Member mainly consists of radiolarian-bearing muddy limestones with calciturbidites.
118 Shale horizons occur in its lower portion. The Middle Streppenosa Member includes

119 mudstones/wackestones, with intraclastic-peloidal and oolitic thin intercalations, often recrystal- lized
120 or dolomitized, and black silty shales. The Upper Streppenosa Member consists of gray-green shales,
121 marls and radiolarian-bearing muddy limestones with calcarenite intercalations. Compared with the
122 Lower and Middle members, the silty shales and quartz siltstones in- crease, while the organic matter
123 decreases (Frixia et al., 2000). Basalts and tuff layers occur in the sedimentary sequence at various
124 strati- graphic levels and become more frequent in the Upper Streppenosa Member. On the basis of
125 previous studies, basaltic horizons have an overall intraplate alkaline nature (Rocchi et al., 1996). The
126 drowning of the carbonate platform-basin system first occurred with the spread of the basinal facies
127 (Upper Streppenosa Member) over the marginal- inner platform complex, as also reported at a
128 regional scale (Catalano et al., 2013). According to previous authors, the drowning phase first oc-
129 curred in the Early Jurassic time (Patacca et al., 1979; Brosse et al., 1988; Frixia et al., 2000) (Fig. 3).

130 - Within this stratigraphic framework, the main elements of the Hyblean Plateau Petroleum
131 System are as follows.

132 - Two reservoirs, represented by the carbonate platform succession (Sciacca Formation) and the
133 marginal microbial mound complex (Mila Member). In the Sciacca Formation porosity reaches
134 maxi- mum values of 20%, mostly due to fracturing (Mattavelli et al., 1969; Frixia et al., 2000).
135 In the Mila Member, the poor reservoir properties are due to prevailing muddy facies but are
136 improved by hydrothermal dolomitization, fracturing and karst.

137 - Two source rock units, represented by the Noto Formation whose Total Organic Carbon (TOC)
138 reaches about 13% and by the Streppenosa Formation. Although the average TOC of the
139 Streppenosa Formation is low (around 1%), its great thickness makes this formation a good
140 source rock (Brosse et al., 1988; Frixia et al., 2000).

141 - One seal, consisting of the upper portion (Upper Streppenosa Mem- ber) of the Streppenosa
142 Formation (Frixia et al., 2000).

143 The investigated wells, Streppenosa 1 and Bimmisca 1, were drilled in the northwestern and eastern
144 parts of the Hyblean Plateau, respectively. The Streppenosa 1 well ($36^{\circ}50'45''\text{N}/02^{\circ}16'09''\text{E}$),
145 reached a depth of 2908.4 m in the upper part of the Sciacca Formation. The Bimmisca 1 well
146 ($36^{\circ}48'34''\text{N}/02^{\circ}37'28''\text{E}$) terminated at a depth of 3169 m within the Mila Member of the Noto
147 Formation, and thus never reached the Sciacca carbonate platform. All the depths were mea-
148 sured below the Rotary Table (MDBRT). The Streppenosa 1 and the Bimmisca 1 wells penetrated the inner
149 area and the marginal complex of the carbonate platform-basin system, respectively (Figs. 1, 2). In
150 this area, only the Upper Streppenosa Member is present, directly over- lying the Noto Formation
151 (including the Mila Member). The complete Streppenosa Formation, including all the three members,
152 is present only in the depocenter of the basin, and is penetrated by the Polpo 1 and Pachino 4 wells
153 (Frixia et al., 2000; Cirilli et al., 2015) (Figs. 2, 3).

154 The succession investigated in the Streppenosa 1 borehole includes the Upper Streppenosa Member
155 (2126 m–2452 m) and the Noto For- mation (2452 m–2825 m). In the Bimmisca 1 well, the
156 succession investigated encompasses the Upper Streppenosa Member (2282 m– 2684 m) and the
157 Noto Formation (2684 m–2803 m), including the Mila Member (2803 m–2879 m). The lowermost
158 portion of the Mila Member, between 3169 m to 2908 m, has not yielded samples due to a loss of
159 circulation. In both wells studied, the Streppenosa Formation is overlain by the pelagic deposits of the
160 lower-middle Jurassic Modica Formation (Patacca et al., 1979), which is not included in this study.

161

162 **3. Methods**

163 Palynological and palynofacies analyses were carried out on a total of 59 cutting samples: 30 from the
164 Streppenosa 1 well and 29 from the Bimmisca 1 well. The samples were processed using standard
165 paly- nological techniques in order to obtain a final organic residue observ- able under a microscope
166 (Green, 2001; Wood et al., 2002; Buratti and Cirilli, 2011). The palynomorphs and palynomacerals

167 were visualized with a Leica DM1000 microscope using transmitted white light. Optical palynofacies
168 analyses were conducted on the organic matter prior to oxidation with nitric acid. The organic
169 particles were classified according to the standard classification of Whitaker (1984), modified by
170 Steffen and Gorin (1993). Marine and terrestrial palynomorphs were identified from both artificially
171 oxidized and unoxidized residue.

172 For this study, only cuttings were available. A common problem with well cuttings is the potential
173 contamination of rock fragments derived from uphole. This could represent a problem when
174 interpreting both the palynostratigraphy and palynofacies. However, in our opinion, using a rigorous
175 approach and method, conclusions can be drawn regarding which palynological assemblages and
176 palynofacies composition can be used as complementary tools. First, to identify the presence of
177 caving contamination, we carefully handpicked the lithologies and compared them with up-hole
178 samples. The identification was made on thin sections made up from cutting material, previously
179 embedded in epoxy resin, observed under transmitted light microscope. Then we used the highest
180 occurrences of taxa to define the tops of the palynozones. The first downhole occurrences (FDOs) of
181 palynomorphs were used in constructing the range charts, in order to minimize the error introduced
182 by caving. The presence of a few anachronistic palynomorphs throughout the studied well succession
183 indicated that contamination from caving was minimal. Thus we attempted to use the palynofacies
184 composition and variation in order to reconstruct the depositional environment.

185 Palynomorphs and all the organic debris were measured quantitatively. At least 250 particles per
186 slide (two slides per sample) were counted and converted into percentages. Details on the sample
187 processing, the general definition of the organic debris and the palynomorph quantitative analysis
188 are given in the Online Resources. The palynological slides were then stored in the collection of the

189 Sedimentary Organic Matter Laboratory at the Department of Physics and Geology of the Perugia
190 University (Italy).

191 **4. Palynological data**

192 A complete list of the identified taxa and data on quantitative analysis can be found in the Online
193 Resource (Appendix I, Figs. S1; S2).

194 *4.1 Palynology of the Streppenosa 1 well*

195 The palynological assemblages in the Streppenosa 1 succession contain abundant sporomorphs in
196 association with minor marine elements (microforaminiferal linings, dinoflagellate cysts and acritarchs)
197 (Fig. 4, Pl.I). The microflora is dominated by *Classopollis meyerianus* and by trilete fern spores, such
198 as *Deltoidospora mesozoica*, *Dictyophyllidites mortonii*, *Todiporites* sp., *Trachysporites fuscus*, which
199 abundances vary through the studied section, in association with minor *Acanthotriletes varius*,
200 *Baculatisporites* sp., *Calamospora tener*, *Carnisporites spiniger*, *Densosporites fissus*,
201 *Kraeuselisporites* sp. and *Porcellispora longdonensis*. The distribution of *Limboisporites lundbladiae*,
202 *Perinopollenites elatoides*, and bisaccate pollen grain as *Klausipollenites gouldii* is not constant
203 throughout the well. The microflora gradually diversify throughout the Upper Streppenosa Member
204 with the first downhole occurrences (FDOs) of *Punctatisporites fungosus*, *Paraklukisporites foraminis*,
205 *Classopollis torosus* and *Striatella seebergensis* in its upper part, and of *Trachysporites fuscus*,
206 *Cingulizonates rhaeticus*, *Densosporites foveocingulatus*, and *Ricciisporites* sp., in its central-lower
207 part. The FDO of the index species *Ischyosporites variegatus* is recorded in the central-upper part
208 (2235.5 m) whereas that of *Porcellispora longdonensis* in the lower portion (2421.5 m) of the member.
209 The presence of *Leptolepidites major*, *Pilosisporites* sp. and *Trilobosporites aequiverrucosus* is
210 considered here as caving. *Ischyosporites variegatus* in association with *Polypodiisporites*
211 *polymicroforatus*, *Retitriletes semimuris*, *Triancoraesporites ancorae*, rare *Araucariacites australis*
212 have their last downhole occurrences (LDO) at the base of the Upper Streppenosa Member (sample at

213 2445.5 m) and are not recorded in the underlying Noto Formation. The assemblage from the Noto
214 Formation is less diversified and records few FDOs in the central-upper part: the FDO of
215 *Eucommiidites* sp. at 2503.5 m and that of *Tsugaepollenites pseudomassulae* at 2555.5 m. The fresh
216 water chlorococcale alga *Botryococcus* sp. is more or less constantly present from the middle part of
217 the Noto Formation slightly decreasing upward, while *Schizosporis scissus* is recorded only at the base
218 of the member. Microforaminiferal linings are constantly present within the investigated well section.
219 Acritarchs and dinoflagellate cysts are rare and commonly poorly preserved. The FDO of *Suessia*
220 *swabiana* is recorded at the base of the Upper Streppenosa Member.

221

222 4.2 Palynology of the Bimmisca 1 well

223 As in the Streppenosa 1 well, the palynological assemblages from Bimmisca 1 well are dominated by
224 sporomorphs in association with minor marine components (Fig. 5, Pl.II). The FDO of *Ischyosporites*
225 *variegatus* is recorded at the top of the studied well section (2302 m) with *Araucariacites australis* and
226 *Deltoidospora mesozoica* and, 19 m below, by *Calamospora tener*. The assemblage diversity and
227 abundance progressively increase downhole with the FDO of *Classopollis meyerianus* at 2481 m. The
228 middle portion of the Upper Streppenosa Member is characterized by the FDO of some important taxa
229 such as *Polypodiisporites polymicroforatus*, *Convolutispora klukiforma*, cf. *Retitriletes semimuris*,
230 *Annulispora folliculosa*, *Limboisporites lundbladiae*, cf. *Retitriletes austroclavatidites*, *Trachysporites*
231 *fuscus*, *Acanthotriletes varius*, *Camarozonosporites rudis*, *Densosporites foveocingulatus*,
232 *Leptolepidites reissingeri*. The FDO of the species index *Porcellispora longdonensis* is recorded at
233 2603 m and that of *Classopollis torosus* at 2610 m.

234 Overall, the fern spores are abundant in the Upper Streppenosa Member and can be found
235 throughout the well section. The alga *Botryococcus* sp. is occasionally present in the middle part of the
236 Upper Streppenosa Member. Microforaminiferal linings, although low abundant, become more

237 constant in the Upper Streppenosa Member. The FDO of *Suessia swabiana* is recorded at the base of
238 the Upper Streppenosa Member, where it is low abundant. The assemblage diversity decreases in the
239 Noto Formation. A few FDOs are recorded in its lower portion as for the spores *Conbaculatisporites*
240 sp., *Conbaculatisporites spinosus* and *Punctatisporites fungosus*.

241

242 **5. Palynofacies and lithofacies data**

243 The use of palynofacies, combined with other approaches, can provide useful information on
244 sedimentary processes and on the chemical and ecological parameters of the depositional environment,
245 such as, oxygenation, length of transportation, water energy and nutrient level (van der Zwan, 1990;
246 Steffen and Gorin, 1993; Hart et al., 1995; Tyson, 1995, 2001; Batten and Stead, 2007). The abundance
247 of total terrestrial organic matter commonly decreases with the distance from the land-masses and the
248 parent flora. More labile components (i.e. cutinite) are rapidly degraded during transport, while
249 oxidized particles (i.e. inertinite) can be widely distributed into marine basins because they are
250 refractory to further biochemical oxidation. In a marine setting, the narrow width of the continental
251 shelf, as expected for the study area, exerts a strong control on the relative positions of proximal and
252 distal environments (Masclé et al., 1996). Important controlling factors on organic matter accumulation
253 and preservation rate include the primary productivity and the oxygen level of the water column.
254 Under suboxic-anoxic conditions, the organic matter is usually converted into amorphous organic
255 matter (AOM). Given that AOM is highly susceptible to oxidation, its presence can be a proxy for the
256 low oxygen content of the water column and sediment water interface (Tyson, 1995; Batten and Stead,
257 2007). The relative abundances of sedimentary organic matter and palynofacies composition from the
258 Streppenosa 1 and the Bimmisca 1 wells are plotted in Figs. 6 and 7. Additional information on the
259 general characteristic of organic debris is provided in the Online Resources.

260

261 *5.1 Streppenosa I well*

262 Six intervals, characterized by different lithofacies and palynofacies, were distinguished in this well.
263 Lithology and stratigraphic distribution of the palynofacies ratios as well as the number and spacing of
264 analyzed samples for each interval are illustrated in Fig. 6.

265 - Interval I (Palynofacies A, Pl. III.1) (2824 m – 2735.5 m) represents the lowest part of the Noto
266 Formation. The interval mainly consists of shales and thin bedded limestones. The palynofacies
267 contains abundant inertinite, which reaches the highest values in the upper part of the interval (58%).
268 Vitrinite 1 abundance decreases from bottom to top (32% to 8%). The amount of vitrinite 2 is almost
269 constant within the interval (7% to 10% upwards). Cutinite, sporomorphs (tetrads included) have a low
270 frequency (less than 5%). Marine components are rare and mainly represented by microforaminiferal
271 linings in the lower part. A few specimens of the freshwater alga *Botryococcus* sp. were recorded in the
272 upper part of the interval. The AOM abundance increases from the base (4%) to the central part (20%)
273 and decreases (15%) at the upper part of the interval .

274 - Interval II (Palynofacies B, Pl. III.2) (2735.5 m – 2624.5 m) consists of limestones intercalated with
275 shales, which become more frequent in the upper part. Vitrinite 1 abundance increases from the lower
276 (38%) to the central part of the interval (58%), and decreases upwards (15%), except for at the interval
277 2637.5. Inertinite and vitrinite 1 abundances show general opposite trends being inertinite lower where
278 vitrinite 1 higher and vice versa. The vitrinite 2 content is overall low, ranging from 2.5% to 12.5%.
279 Sporomorphs, tetrads included, relatively increase in the central part of the interval (up to 15%),
280 cutinite amount is low (less than 2%) within the interval. Microforaminiferal linings and *Botryococcus*
281 sp. slightly increase upwards (up to 5%). The highest AOM values (up to 30%) are recorded in the
282 shaly beds at the lower part of the interval.

283 - Interval III (Palynofacies C, Pl. III.3) (2555.5 m – 2476.5 m), overlying a basalt horizon, is composed
284 of thick shaly horizons alternated to limestone. Inertinite and vitrinite 1 fluctuate along the interval with

285 a peak of respectively 54% and 45%, showing an opposite trend. Vitrinite 2 abundance ranges between
286 8% to 16%. The amount of cutinite slightly increases in respect to the underlying interval (up 6%), that
287 of sporomorphs slightly decreases and tetrads are few. The AOM reaches the maximum value (30%)
288 within the shaly intervals in correspondence of the lowest value of vitrinite 1 and moderate amount of
289 inertinite. Microforaminiferal linings are few. A few specimens of *Botryococcus* sp. are recorded only
290 in the lower and central part of the interval.

291 - Interval IV (Palynofacies D, Pl. III.4) (2476.5 m – 2409.5 m) includes the Noto Formation and Upper
292 Streppenosa Member boundary (at 2452.5 m depth). The interval exhibits a basal thick carbonate
293 horizon (Noto Formation) passing upwards into alternations of dark grey shales and limestones
294 containing fine grained calcarenites (Upper Streppenosa Member). The palynofacies shows the highest
295 values of AOM (up to 40%) at the lower and at the upper part of the interval, the latter is combined
296 with high values of pyrite framboids. In the central part of the interval, AOM abundance decreases up
297 to 5%. There is a progressive upward increase in vitrinite 2 (10% to 35%) accompanied by a decrease
298 in vitrinite 1 (28% to 6%), while inertinite values remain almost constant. The sporomorph abundance,
299 tetrads included, shows a peak in the central part of the interval (up to 25%). Cutinite amount fluctuates
300 from low values (1.5%) in the central part and relative higher values at the lower and upper part of the
301 interval. Constant low percentage (5%) of marine elements (microforaminiferal linings and few
302 dinoflagellate cysts) has been recorded in the central and upper part of the interval. Low amount of
303 *Botryococcus* sp. is present at the base and at the top of the interval.

304 - Interval V (Palynofacies E, Pl. III.5) (2409.5 m – 2308.5 m) consists of black shales in the lower part
305 passing, upwards, to thick carbonate beds, which contain calcarenites, interbedded with shaly intervals
306 and radiolarian bearing muddy limestones. The low number of samples prevented a detailed
307 palynofacies characterization of this interval that has been possible only from 2357.5 m to 2308.5 m.
308 Inertinite reaches the highest values (55%) at the base of the sampled interval, where all the other

309 palynomacerals and sporomorphs decrease. Vitrinite 1 slightly increases upwards (up to 25%), while
310 vitrinite 2 abundance remains constant (15%) as well as cutinite and sporomorphs although occurring
311 in low abundance. The AOM values increase in the upper part of the interval (25%) within the shaly
312 interval. A few microforaminiferal linings (1.8%) are present.

313 - Interval VI (Palynofacies F, Pl. III.6; IV.1) (2308.5 m – 2147.5 m) is dominantly carbonate except for
314 a basal thick dark shaly horizon and thin intercalations of dark shales within carbonates. The carbonate
315 horizons consist of thin bedded limestone mostly characterized by radiolarian and thin shelled bivalve
316 bearing mudstone-wackestone. Due to the lack of samples, the palynofacies analysis has been possible
317 starting from the central part of this interval. Vitrinite 1 and sporomorphs record the highest value in
318 the central part of the interval (58% and 22% respectively). Vitrinite 2 abundance fluctuates within the
319 interval (8% to 20%) as well as cutinite (5% to 10%). The AOM shows the highest value in the central
320 part of the interval (40%), whereas pyrite is present only in the central (16%) and upper (6%) part. The
321 percentage of microforaminiferal linings increases upwards and *Botryococcus* sp. is consistently
322 present (10%) at the topmost interval.

323

324 5.2 *Bimmisca* 1 well

325 Five intervals and related lithofacies and palynofacies were recognized. Lithology and stratigraphic
326 distribution of the palynofacies ratios, the number and spacing of analyzed samples are illustrated in
327 Fig. 7.

328 - Interval I (Palynofacies G, Pl. IV.2) (2879 m – 2823 m), in the Mila Member, composed mainly of
329 microbial dolostones. Terrestrial organic debris dominates the palynofacies. Vitrinite 1 reaches high
330 values (73%) in the central-upper part of the interval, whereas vitrinite 2 ranks at low percentages
331 (around 5%). Few sporomorphs are present (1.5%). Cutinite abundance shows moderate value (19%) at
332 the base of the interval and decreases upwards. Inertinite and AOM values show the same pattern

333 progressively decreasing upwards (40% to 5%, inertinite; 38% to 15%, AOM). Microforaminiferal
334 linings and acritarchs are few (1%).

335 - Interval II (Palynofacies H, Pl. IV.3) (2823 m – 2694 m) mostly belongs to the Noto Formation
336 overlying the Mila Member. It consists of dolostones intercalated with minor shales. Vitrinite 1 and
337 inertinite fluctuate (10% up to 80%) within the interval showing an opposite trend. Vitrinite 2
338 abundance is almost constant ranging from 3% to 7%. Similarly the amount of cutinite remains more or
339 less constant within the interval. Sporomorph abundances (tetrad included) show variations along the
340 interval never exceeding the 18%. The AOM values are variable (25% to 2%) showing an overall
341 decreasing trend upward. Microforaminiferal linings and acritarchs are few (up to 2%).

342 - Interval III (Palynofacies I, Pl. IV.4) (2694 m – 2603 m). The base of this interval coincides with the
343 Noto Formation - Upper Streppenosa Member boundary. The interval is marked by an abrupt decrease
344 in carbonate content, being dominated by shales with thin muddy carbonate intercalations. Overall, the
345 palynofacies shows an increase in the total amount of continental organic debris and a decrease in
346 AOM (from 20% to 2% upwards). Inertinite and vitrinite 1 show strong fluctuations, the former
347 reaching the highest values in the carbonate beds. Vitrinite 2 ranks at low abundance (2% to 10%) as
348 well as cutinite. Sporomorphs and tetrads increase upwards ranking at 32%, the highest value recorded
349 within the well section. Marine palynomorphs (microforaminiferal linings, acritarchs and few
350 dinoflagellate cysts) slightly increase in respect to the underlying intervals.

351 - Interval IV (Palynofacies L, Pl. IV.5) (2603 m – 2441 m) consists of shales with minor intercalations
352 of carbonate composed of thin bedded muddy limestone. The palynofacies is dominated by inertinite
353 (up to 92% in the upper part of the interval). The other palynomacerals and sporomorphs are low to
354 absent, except for moderate value of vitrinite 1 (up to 30%). AOM content is low (less than 8%) and
355 further decreases to disappear upwards. Pyrite (8%), marine components (not more than 5%) such as

356 microforaminiferal linings and rare acritarchs are present in the lower part. *Botryococcus* sp. has been
357 recorded only in this interval.

358 - Interval V (Palynofacies M, Pl. IV.6) (2342 m – 2302 m.) overlies a basalt horizon. It consists of
359 shale and thin bedded muddy limestone alternations. Palynofacies is dominated by inertinite (45% to
360 84%) and vitrinite 1 (up to 45%), combined with low amount of other palynomacerals and
361 sporomorphs. Tetrads are absent. The abundance of AOM is low (up to 6%) as well as that of pyrite
362 (up to 2%), this latter found only in the upper part of the interval. A few microforaminiferal linings and
363 acritarchs are present (up to 4%).

364

365 **6. Discussion**

366 *6.1 Palynostratigraphic assessment and dating*

367 The Noto Formation and the Upper Streppenosa Member can be assigned to the Rhaetian age given the
368 abundance of *Classopollis* species (*Classopollis meyerianus* and minor *Classopollis torosus*) in
369 assemblage with other index species such as *Ischyosporites variegatus*, *Porcellispora longdonensis* and
370 *Trachysporites fuscus*. These species have been found in typical Rhaetian strata from several localities
371 (Morbey, 1975; Schuurman, 1977; Barrón et al., 2006; Kürschner et al., 2007; Warrington et al., 2008;
372 Ruhl et al., 2009; Cirilli, 2010; de Jersey and McKellar, 2013; Hillebrant et al., 2013; Lindström, 2016;
373 Lindström et al., 2017b). Palynological assemblages with similar compositions have been recorded
374 from Southern Alps and Apennines in Italy (Cirilli et al., 1994; Galli et al., 2007) and in some southern
375 Mediterranean areas such as Tunisia, Libya, Algeria, Morocco (Adloff et al., 1986; Yaroshenko, 2007;
376 Cirilli, 2010). Palynological assemblages from Northern Spain (Asturias) and France, although
377 showing some common elements (e.g. dominance of *Classopollis* spp. and fern spores) differ for the
378 presence of *Ricciisporites tuberculatus* (Barrón et al., 2006; Gómez et al, 2007). The absence of *R.*
379 *tuberculatus* could be related to the palaeogeographic position of Sicily during Late Triassic along the

380 African- European plate boundary, under warm paleoclimatic conditions (Ruiz-Martínez et al., 2012;
381 Catalano et al., 2013; Berra and Angiolini, 2014; Scotese and Schettino, 2017). Recent data have
382 documented that the gymnosperm pollen *R. tuberculatus* appears to be more restricted to the Northern
383 Hemisphere (Kürschner et al., 2014; Lindström, 2016; Lindström et al., 2017a). Its abundance seems to
384 be vicariant with that of *Classopollis*, being less common or absent where *Classopollis* is abundant, as
385 in the present case, revealing different ecological and/or climatic preferences. In the GSSP stratotype
386 Kuhjoch section (Karwendel Mountains, Austria) the lowest occurrences of *I. variegatus* and
387 *Cerebropollenites thiergartii*, which is considered the best marker for the T-J boundary, are recorded
388 several meters below the first appearance of the ammonite *Psiloceras spelae* defining the base of the
389 Hettangian (Bonis et al. 2009; Kürschner and Hengreen, 2010; Hillebrant et al., 2013). Therefore, the
390 presence of *I. variegatus* without *C. thiergartii* and in association with *Classopollis* spp., *P.*
391 *longdonensis* and *Kraeuselisporites reissingeri*, led tentatively to refer part of the Upper Streppenosa
392 Member to the lowermost part of the *Trachysporites–Heliosporites* Zone (TH) which is considered as
393 Rhaetian (Hillebrant et al., 2013). The thickness of the interval below the FO of the *C. thiergartii*
394 depends from the sedimentation rate which surely was not the same for all the sections bracketing the
395 T/J boundary. In this case study, considering the high subsidence rate caused by synsedimentary
396 tectonic activity and the medium-to high sedimentation rate (Patacca et al., 1979), the presence of *C.*
397 *thiergartii* in the upper part of the Upper Streppenosa Member cannot be excluded. However its
398 presence has not been recorded either in these two wells or in the Pachino 4 well (Cirilli et al., 2015).
399 The microfloral content recorded in the lower part of the Upper Streppenosa Member and in the Noto
400 Formation dominated by *Classopollis meyerianus* in association with *T. fuscus*, *P. longdonensis*,
401 *Kraeuselisporites* sp. and *Polypodiisporites polymicroforatus* could be correlated with the
402 *Trachysporites–Porcellispora* Zone (TPo) considered as latest Rhaetian as defined in the Tiefengraben
403 section and Northern Calcareous Alps, Austria, (Kürschner et al. 2007; Hildebrandt, 2013). The age

404 attribution of the whole studied well section to latest Rhaetian would explain also the lack of
405 *Patinasporites densus* and *Enzonalasporites vigens* which seems to have their last occurrences in the
406 uppermost Norian-lower Rhaetian (Cirilli, 2010 for references). In Sicily these two species have been
407 recorded in independently dated Carnian and Norian strata both from outcrops and subsurface
408 (Visscher and Krystyn, 1978; Buratti and Carrillat, 2002; Trincianti et al., 2015).

409 By correlating the new palynological data with those of Pachino 4 well (Cirilli et al., 2015), it results
410 that the deposition of Noto Formation, at the marginal and inner carbonate platform-basin system, may
411 be considered coeval with the Middle Streppenosa Member deposited in the deepest part of the basin.
412 Furthermore, it implies that the initial drowning phase of the carbonate-basin system (Upper
413 Streppenosa Member) can be predated to Rhaetian. The previous age attribution as Early Jurassic of the
414 whole Upper Streppenosa Member (Fig. 3) was based on the presence of the calcareous nannofossil
415 *Schizosphaerella punctulata* and on a palynological assemblage dominated by *Classopollis classoides*
416 and *C. meyerianus* and (Frixia et al., 2000). However, although *S. punctulata* was most common in the
417 Early Jurassic times its range spans from the latest Triassic to the end of the Jurassic (Perch-Nielsen,
418 1989). In St Audrie's Bay (England), *S. punctulata* first occurs about one metre below the candidate
419 Hettangian GSSP level (in Blue Lias Formation bed 7). Its occurrence slightly predates the base of the
420 *Psiloceras planorbis* Zone and correlates with the onset of the main negative excursion in $\delta^{13}\text{C}_{\text{org}}$
421 values (Hesselbo et al., 2002, 2004). Additionally, the relative abundance of *Classopollis* spp., which is
422 also present in the Triassic, does not justify assigning a Jurassic age to the Upper Streppenosa Mem-
423 ber, given that in this case *Classopollis* spp. do not occur alongside typical Jurassic forms.

424

425 6. 2 Palaeoenvironment, palaeoclimate and depositional model

426 Considering the general paleogeographic and paleoenvironmental settings, the lateral and vertical
427 variations of palynofacies and lithofacies and the prevailing continental organic matter across the well

428 sections indicate a deposition in a marine environment, proximal to the terres-
429 variable energy and redox conditions.

430 The Noto Formation, crossed by the Streppenosa 1 well, located in the inner part of the carbonate
431 platform-basin system, has a clear cyclic arrangement. Each interval shows a trend characterized by a
432 decreasing AOM and palynomaceral and sporomorph increasing upwards. The rel- ative abundance of
433 AOM in the basal part of each interval results from a combination of good preservation, related to
434 suboxic–anoxic condi- tions, and low-energy environments. It has also resulted in the deposi- tion of
435 laminated shales with few limestone intercalations. The upper part of each interval shows shallower,
436 and relatively better oxygenated, conditions where particulate organic matter dominates over AOM,
437 which was presumably destroyed by oxidation and biodegradation. In the Bimmisca 1 well, which
438 penetrates the platform marginal area, the palynofacies fluctuations are more evident at the scale of the
439 entire suc- cession (Noto Formation/Mila Member and Upper Streppenosa Mem- ber), in accordance
440 with lithofacies variation from mainly carbonate to predominantly shaly upwards. The increase in
441 inertinite and the de- crease in terrestrial particulate organic matter and AOM from bottom to top are
442 consistent with a shift from the proximal to distal position of the platform-basin system with respect to
443 the landmasses. The moderate-to-high value of AOM recorded in the shallow water micro- bial mounds
444 of the Mila Member could be partially related to the micro- bial community itself. Under microbial
445 proliferation, the extracellular polymeric substances (EPS) form a well–defined protective envelope
446 around cyanobacteria cells, which protect AOM from the oxidation pro- cesses (Sutherland, 2001;
447 Pacton et al., 2007). The overall deepening trend of the platform-basin system culminated with the
448 deposition of the Upper Streppenosa Member. In each interval, palynofacies signa- tures clearly reflect
449 the lithofacies variations, which consist of thick shaly beds and minor limestones containing calcarenite
450 intercalations: dominant AOM and minor inertinite in the shaly intervals, highest amount of inertinite
451 and lowest amount of AOM within limestones. These fluctuations could be interpreted as a result of

452 changes in water energy and oxygen content. The higher energy and good oxygenated conditions
453 during the calciturbidites deposition destroyed the AOM and reduced the preservation rate of
454 particulate OM, except inertinite, which is the most resistant palynomaceral. The scarce benthic fauna
455 living at such deep marine bottoms may also have played a role. On the other hand, the suboxic-anoxic
456 conditions, during the deposition of finely laminated shales, promoted the AOM preservation. The
457 concu- rent AOM decrease and inertinite increase upwards confirm the gradual and relative deepening
458 of the depositional environment, highlighted by the basinal facies of the Upper Streppenosa Member
459 onlapping and overlying the microbial mound (Mila Member) at the margin complex and the inner
460 suboxic lagoonal facies of the Noto Formation. The increased deepening and the facies onlapping have
461 been widely reported at the regional scale (Catalano et al., 2013). The drowning phases of the
462 carbonate platform-basin system were accompanied by depositional environment shifting from
463 proximal to distal conditions with respect to the landmasses and parent flora. This is highlighted in both
464 wells by the palynofacies signatures, and the decrease, upwards, of the total terrestrial organic debris
465 (i.e. vitrinite, cutinite and sporomorphs, tet- rads included) with the exception of inertinite. Given that
466 inertinite is the most resistant palynomaceral, it can be transported for a long dis- tance from the
467 continent and settled from suspension in a low-energy, offshore environment (Steffen and Gorin,
468 1993). Based on the data ob- tained and correlated with data from the Pachino 4 well, belonging to the
469 same petroleum system and discussed in a previous paper (Cirilli et al., 2015), the paleoenvironmental
470 evolution of the entire platform- basin system could be interpreted in terms of an “expanding puddle
471 model” (sensu Wignall, 1991; Wignall and Newton, 2001). This model suggests the development of
472 anaerobic conditions within relative shal- low intra-cratonic confined basins, with deposition of organic
473 rich sed- iments both in the deepest part of the basin (i.e. Pachino 4 well, Cirilli et al., 2015) and in the
474 shallow marine areas close to the margin of the carbonate platform (crossed by the Streppenosa 1 and
475 the Bimmisca 1 wells) colonized by microbial mounds (Mila Member) (Fig. 2). During the Rhaetian, as

476 marine transgression continued, the inner peritidal area of the carbonate platform started to drown
477 under permanent subtidal water conditions, with episodes of low oxygenation (Noto For-
478 mation). However, the carbonate factory continued to produce, as dem-
479 onstrated by the calcarenite intercalations within the organic rich facies deposited in the adjacent basin. The platform margin
480 setting facilitated the development of toe-of-slope aprons characterized by calciturbidites, intercalated
481 with organic rich shales. At the same time, as observed in the Pachino 4 well (Cirilli et al., 2015), in the
482 deepest part of the basin, thick organic rich shales and limestones sedimented. Subsequently, the
483 maximum rise in sea level, combined with the increase in subsi-
484 dence caused the expansion of deep waters and the spread of organic rich facies over the marginal-inner platform complex. This step corre-
485 sponds to the deposition of the Upper Streppenosa Member in the basin depocenter (cf. Pachino well in
486 Cirilli et al., 2015) until onlapping the marginal and inner part of the preexisting carbonate platform.
487 According to our palynological data, the initial drowning of the carbonate-platform can be dated as
488 Rhaetian. Intense tectonic activity related to the continental rifting (and on a larger scale to the Pangea
489 fragmentation) on the northern edge of the African craton (Ruiz-
490 Martínez et al., 2012; Catalano et al., 2013; Berra and Angiolini, 2014; Scotese and Schettino, 2017) was the main cause of the sea level
491 varia-
492 tions in the western Tethys area. In the meantime increasing humidity led to increased freshwater
493 input in the sedimentary basin. This is suggested by the combined presence of fern spores and
494 chlorococcale algae (*Botryococcus*) whose occurrence in marine deposits tends to indicate freshwater
495 incursions (Batten and Grenfell, 1996). This is also demon-
496 strated by the abrupt increase in clay content, found in the middle-upper portion of the well sections. The main cause for this abrupt cli-
497 mate
498 change could be due to the degassing of basalt flows from the CAMP (van de Schootbrugge et al.,
499 2009; Bonis et al., 2010; Schoene et al., 2010; Ruhl et al., 2011; Schaller et al., 2011; Lindström et al.,
2012, 2017b; Pálffy and Zajzon, 2012; Bond and Wignall, 2014; Fijałkowska-Mader, 2015; Davies
et al., 2017). The increasing atmospheric CO₂ concentrations created up to 3–4° in warming, thus

500 leading to a substantial increase in atmospheric water vapor. Consequently the huge amount of
501 greenhouse-gas emissions in the atmosphere and in the ocean waters could be the main cause of the
502 mass extinction at the end of Triassic. Recent data (Davies et al., 2017; Lindström et al., 2017b) date
503 the onset of the CAMP magmatic activity to about 1000kyr before the earliest known eruptions,
504 showing clear evidence of a strict relation between CAMP, increasing humidity and ETE. The
505 location of Sicily in the northern area of the African Plate during the Late Triassic, suggests that the
506 effects of CAMP degassing could have also affected this area.

507 **7. Conclusions**

508 The results of the integrated palynostratigraphy, palynofacies and lithofacies data from the studied well
509 sections shed new light on the Early Mesozoic evolution of the Hyblean Plateau and on the factors trig-
510 gering the organic matter accumulation and preservation in these kinds of intraplatform basins. The
511 palynological assemblages enable the age of the source and seal rocks to be defined and to characterize
512 the timing of the drowning phases of the carbonate platform-basin system. Based on the new
513 palynological data, the entire succession (Noto Formation and Upper Streppenosa Member), can be
514 dated as Rhaetian, thus constraining the initial drowning phase of the carbonate-basin system along a
515 time interval characterized by increasing global humidity, related to the degassing of basalt flows
516 from the CAMP. Climate-driven fluctuations in continental runoff controlled anoxia and black shale
517 deposition in the Triassic Hyblean basin. Humid phases characterized by high precipitation, strong
518 chemical weathering, and freshwater runoff from landmasses caused water stratification and
519 subsequent anoxia at marine bottoms, facilitating the preservation of sedimentary organic matter. The
520 humid phase is highlighted by the increase in clay content and by the presence of fern spores and
521 *Botryococcus* algae. The integrated palynofacies and lithofacies data highlighted a paleogeographic
522 scenario, consisting of a platform-basin system whose evolution was strongly controlled by relative
523 sea level changes triggered by a combination of tectonic and climate factors. Both palynofacies and

524 lithofacies patterns clearly reflect a meter- scale cyclicity in the succession. The paleoenvironmental
525 history of the Hyblean area was strongly marked by a progressive sea-level rise during the Upper
526 Triassic. This is highlighted by the vertical arrangement of cy- cles showing a deepening-upward trend,
527 which culminated with the drowning of the carbonate platform and the spread of organic rich facies
528 over the marginal-inner platform, at the end of the Triassic. This study demonstrates how palynofacies
529 analysis can be used as important com- plementary tool in determining the depositional environment
530 and to de- cipher paleoenvironmental and paleoclimatic changes.

531 Supplementary data to this article can be found online at

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533

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542

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805

806

807 **Figure captions**

808 **Fig. 1** Location of the study wells in the Late Triassic palaeogeographic scenario of the Hyblean
809 Plateau (SE Sicily), at the initial drowing phase (modified from Cirilli et al., 2015); stars indicate the
810 studied wells: Streppenosa 1 and Bimmisca 1, subject of this paper. Circles indicate the location of
811 Pachino 4 well (Cirilli et al., 2015) and Polpo 1 (Frixia et al., 2000).

812

813 **Fig. 2** Palaeoenvironmental restoration and facies distribution during the Late Triassic times of the
814 Hyblean Plateau (not to scale). See the text for explanation.

815

816 **Fig. 3** Scheme of the stratigraphic relationships of the Lower Mesozoic succession and age attribution
817 according to various authors (not to scale). The vertical lines represent the stratigraphic interval
818 covered by the two investigated wells: Bimmisca 1 (Bim1); Streppenosa 1 (Strep1).

819

820 **Fig. 4** Range chart distribution of the terrestrial and marine palynomorphs across the Noto Formation
821 and the Upper Streppenosa Member in the Streppenosa 1 well. Because samples are cuttings, the first
822 downhole occurrences (FDOs) of palynomorphs have been used in constructing the range charts, in
823 order to minimize the error introduced by caving.

824

825 **Fig. 5** Range chart distribution of the terrestrial and marine palynomorphs across the Mila Member,
 826 Noto Formation and Upper Streppenosa Member in the Bimmisca 1 well. Because samples are
 827 cuttings, the first downhole occurrences (FDOs) of palynomorphs have been used in constructing the
 828 range charts, in order to minimize the error introduced by caving. Due to lost of circulation, the
 829 lowermost portion of Mila Member, from 3169 m (Total Depth) to 2908 m, lacks of samples.

830

831 **Fig. 6** Streppenosa 1 well: lithostratigraphic log, lithofacies and palynofacies intervals (I - VI) defined
 832 on the basis of different organic debris percentages. PM1: vitrinite 1 and PM2: vitrinite 2; PM3:
 833 cutinite, PM4: inertinite; SP: sporomorphs (land spores and dispersed pollen grains); T: sporomorph
 834 tetrads; MC: marine components (dinoflagellate cysts, acritarchs and microforaminiferal linings);
 835 Botryoc: *Botryococcus* sp.; AOM: amorphous organic matter.

836

837 **Fig. 7** Bimmisca 1 well: lithostratigraphic log, lithofacies and palynofacies intervals (I - V) defined on
 838 the basis of different organic debris percentages. PM1: vitrinite 1 and PM2: vitrinite 2; PM3: cutinite,
 839 PM4: inertinite; SP: sporomorphs (land spores and dispersed pollen grains); T: sporomorph tetrads;
 840 MC: marine components (dinoflagellate cysts, acritarchs and microforaminiferal linings); Botryoc:
 841 *Botryococcus* sp.; AOM: amorphous organic matter.

842

843 **Plate I** – Palynomorphs from Streppenosa 1 well: 1) *Baculatisporites* sp., Str 28, E.F. O46; 2)
 844 *Perinopollenites elatoides*, Str 28, E.F. J43/1; 3) *Deltoidospora mesozoica*, Str 12, E.F. P38/3; 4)
 845 *Dictyophyllidites mortonii*, Str 21, E.F. J42; 5) *Trachysporites fuscus*, Str 28, E.F. P48/1; 6)
 846 *Porcellispora longdonensis*, STR28, E.F. N29/3; 7) *Tsugaepollenites pseudomassulae*, Str 7, E.F. H36;
 847 8) *Classopollis meyerianus*, Str 28, E.F. N41/3; 9) *Calamospora tener*, Str 29, E.F. F46; 10)
 848 *Carnisporites spiniger*, Str 28, E.F. S36/4; 11) *Acanthotriletes varius*, Str 28, E.F. W45/3; 12) *P.*

849 *polymicroforatus*, Str 28, E.F. N48/3; 13) *Araucariacites australis*, Str 28, E.F. F49/3; 14)
 850 *Ischyosporites variegatus*, Str 30, E.F. B38; 15) *Ischyosporites variegatus*, Str 28, E.F. J46/1; 16)
 851 *Triancoraesporites ancorae*, Str 28, E.F. R34; 17) *Schizosporis scissus*, Str 28, E.F. G39/4; 18)
 852 *Striatella seebergensis*, Str 45, E.F. N45/ 4; 19) *Classopollis torosus*, Str 45, E.F. U52/1; 20)
 853 *Classopollis torosus*, Str 45, E.F. Q24. Scale bar 10 µm.

854

855 **Plate II.** - Palynomorphs from Bimmisca 1 well: 1) *Dictyophyllidites mortonii*, Bim 16, E.F. P38; 2)
 856 *Trachysporites fuscus*, Bim 18, E.F. E51/4; 3) *Trachysporites fuscus*, Bim 17, E.F. T27/3; 4)
 857 *Deltoidospora mesozoica*, Bim 16, E.F. V44/4; 5) *Converrucosisporites* sp., Bim 25, E.F. Q34; 6) *P.*
 858 *polymicroforatus*, Bim 15, E.F. G49/3; 7) *Classopollis meyerianus*, Bim 16, E.F. G32; 8)
 859 *Kraeuselisporites* sp., Bim 16, E.F. N33/4; 9) *Paraklukisporites foraminis*, Bim 18, E.F. F45/1; 10)
 860 *Limbosporites lundbladiae*, Bim 17, E.F. V45; 11) *Verrucosisporites* sp., Bim 17, E.F. P40/1; 12)
 861 *Granulatisporites* sp., Bim 19, E.F. F44/2; 13) *Schizosporis scissus*, Bim 19, E.F. G35/3; 14)
 862 *Perinopollenites elatoides*, Bim 18, E.F. H50/1; 15) *Leptolepidites reissingeri*, Bim 18, E.F. J42/2; 16)
 863 *Porcellispora longdonensis*, Bim 16, E.F. M42/1; 17) *Porcellispora longdonensis*, Bim 16, E.F. E.F.
 864 N42/1; 18) *Ischyosporites variegatus*, Bim 1, E.F. E.F. R40/1; 19) *Retitriletes austroclavadites*, Bim
 865 17, E.F. T50; 20) *Convolutispora klukiforma*, Bim 16, E.F. V39/2. Scale bar 10 µm.

866

867 **Plate III** Palynofacies from the studied interval of Streppenosa 1 well: 1) Interval I-Palynofacies A
 868 (Noto Formation), dominated by inertinite and low percentage of other palynomacerals (Str9); 2)
 869 Interval II-Palynofacies B (Noto Formation) from the shaly intervals, with moderate to high
 870 percentages of inertinite and vitrinite (Str11); 3) Interval III - Palynofacies C (Noto Formation) with
 871 low percentages of vitrinite 2, moderate inertinite and low AOM content from marly limestone (Str23);
 872 4) Interval IV-Palynofacies D (include the Noto Formation and Upper Streppenosa Member boundary),

873 with moderate to high amount of vitrinite, inertinite, sporomorphs and AOM (Str28); 5) Interval V -
874 Palynofacies E, with abundant inertinite and less vitrinite and AOM (Str30); 6) Interval VI -
875 Palynofacies F (Upper Streppenosa Member) with vitrinite, flakes of AOM and less inertinite at the
876 base of the interval (Str36). Scale bar 200 μm .

877

878 **Plate. IV** Palynofacies from the studied interval of Streppenosa 1 well (1) and Bimmisca 1 well (2-6).

879 1) Streppenosa 1 well, Interval VI - Palynofacies F (Upper Streppenosa Member) with abundant AOM,
880 inertinite and less pyrite (Str45); 2) Bimmisca 1 well, interval I - Palynofacies G (Mila Member of the
881 Noto Formation) high content of vitrinite 1 and minor inertinite, moderate AOM from the microbial
882 carbonates at the top of the interval; oil drops are visible (Bim37); 3) Interval II - Palynofacies H (Noto
883 Formation), moderate to high content of inertinite, vitrinite and minor AOM (Bim26); 4) Interval III -
884 Palynofacies I (Upper Streppenosa Member) abundant vitrinite, subordinate inertinite and flakes of
885 AOM in the shaly intervals (Bim24); 5) Interval IV- Palynofacies L (Upper Streppenosa Member)
886 dominated by inertinite with few others palynomacerals (e.g. vitrinite) (Bim14); 6) Interval V -
887 Palynofacies M, shows an abrupt decrease in the total OM content mostly composed of inertinite
888 (Bim1). Scale bar 200 μm .

889

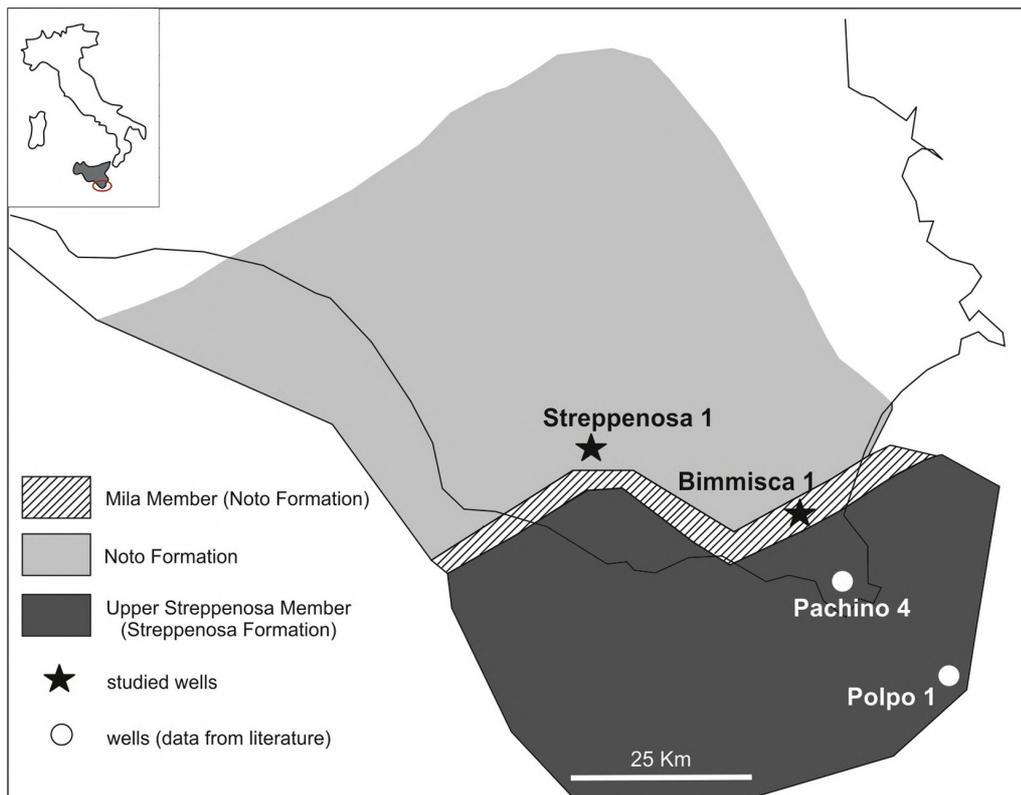


Fig. 1

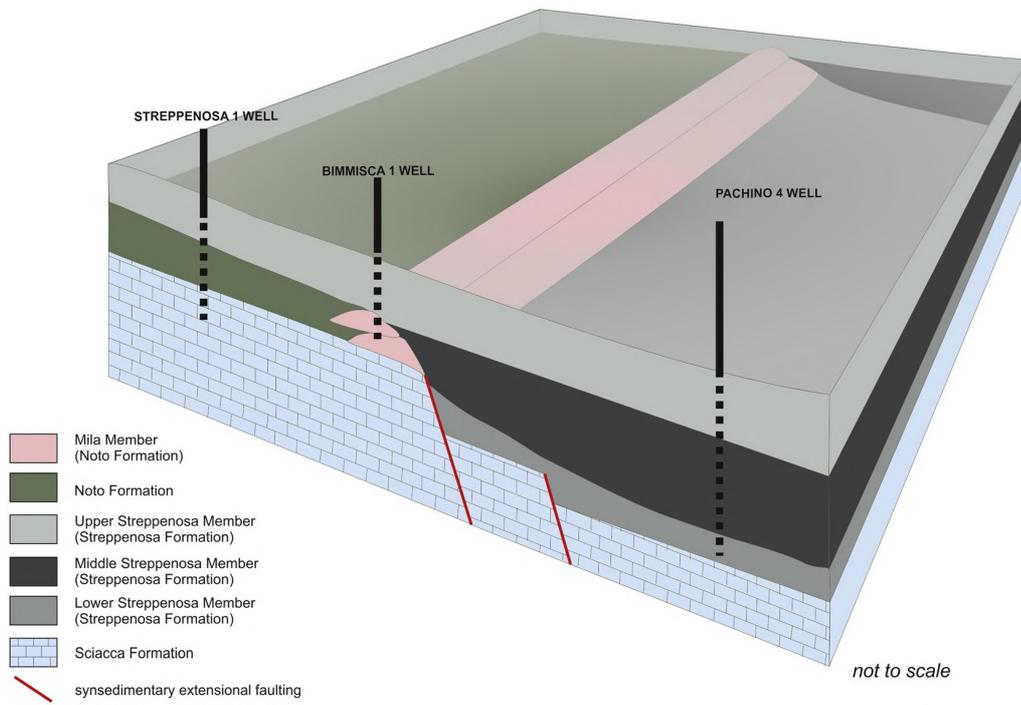


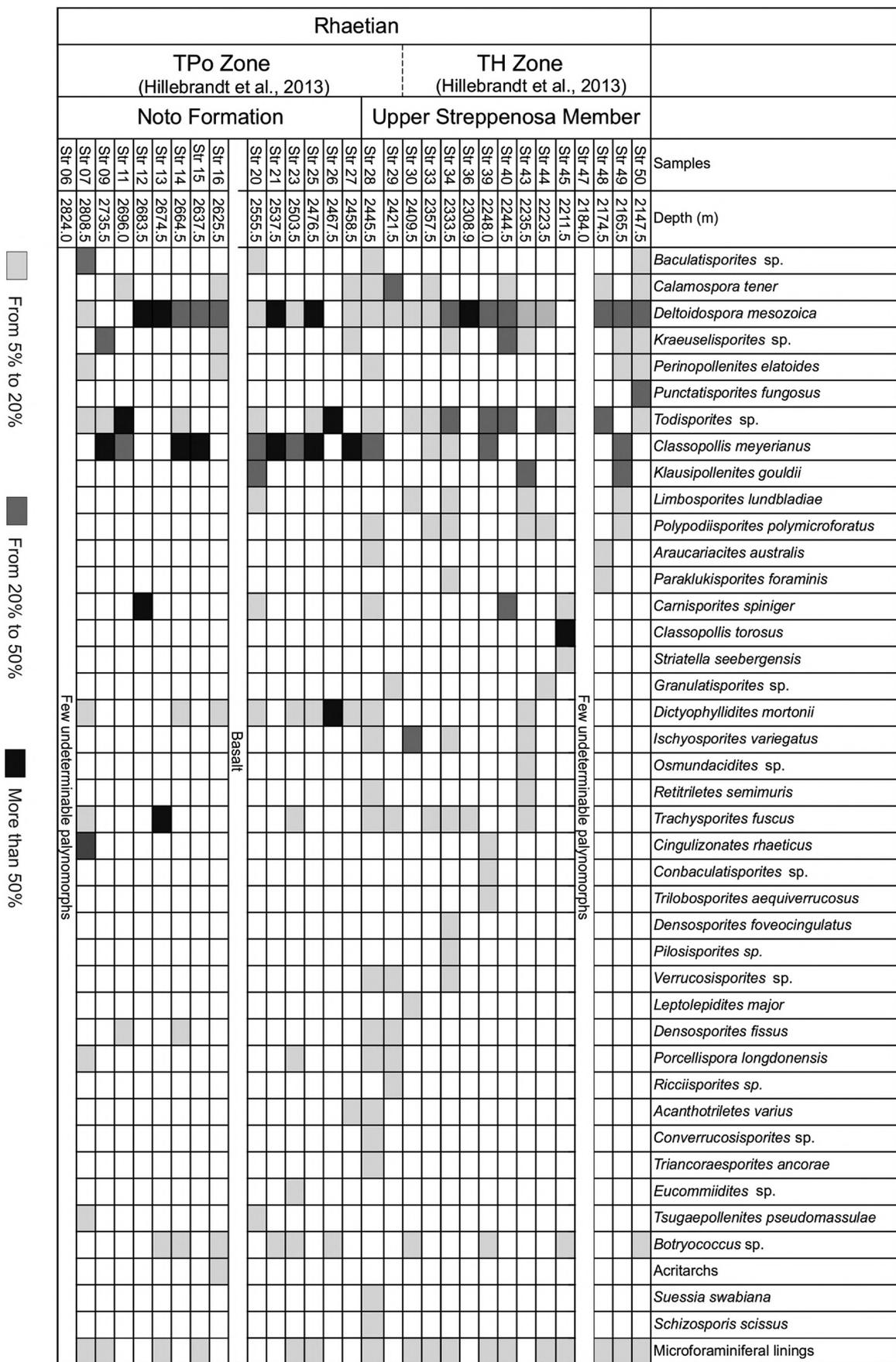
Fig. 2.

	<i>Patacca et al. 1979</i> <i>Brosse et al. 1988</i>	<i>Frixa et al. 2000</i>	<i>Cirilli et al. 2015</i> Basin	<i>This study</i> Inner-marginal complex
Hettangian	Streppenosa Fm.	Upper Streppenosa Mb.		
Rhaetian	Noto Fm.	Noto Fm. (Mila Mb. included)	Upper Streppenosa Mb.	Strep. 1 Upper Streppenosa Mb. Bim. 1
Norian		Lower Streppenosa Mb.	Middle Streppenosa Mb.	Noto Fm. Mila Mb.

Not to scale

Fig. 3.

Fig. 4.



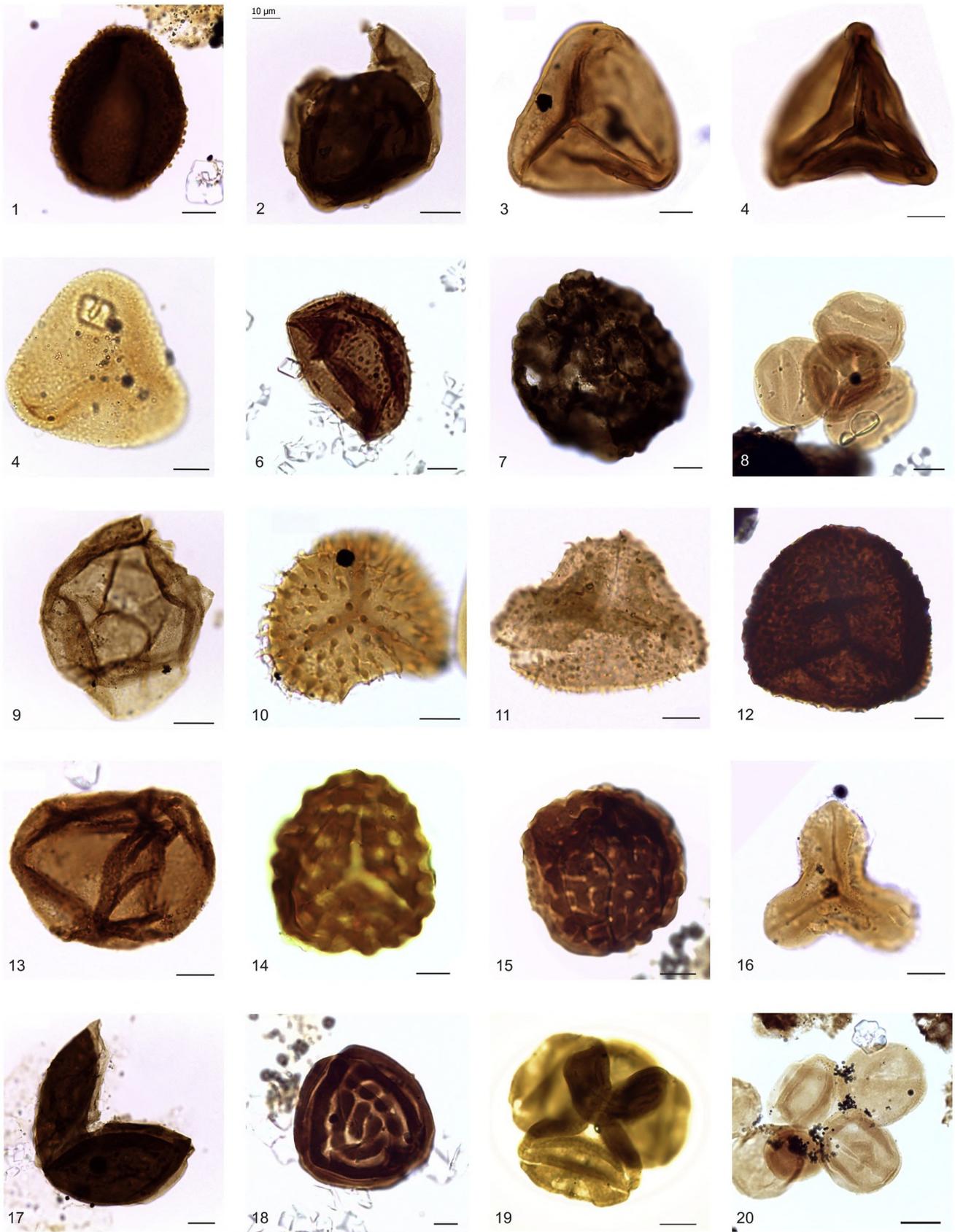
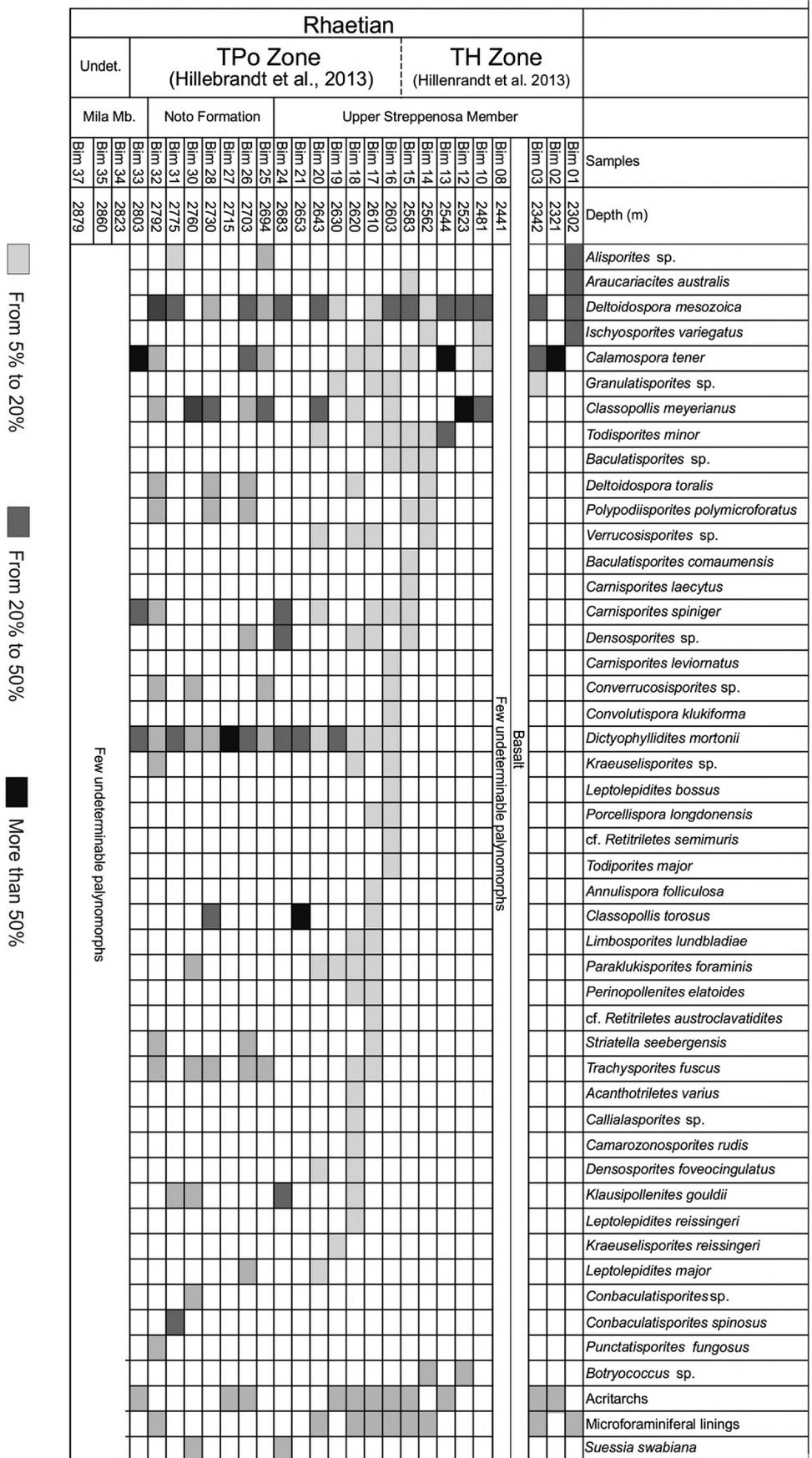
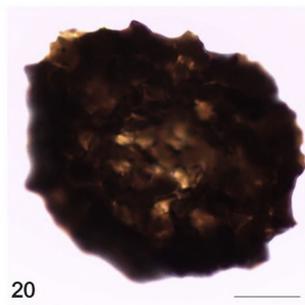
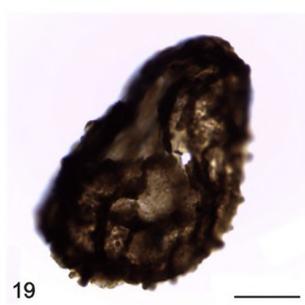
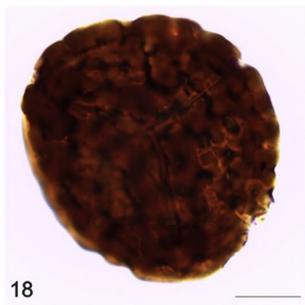
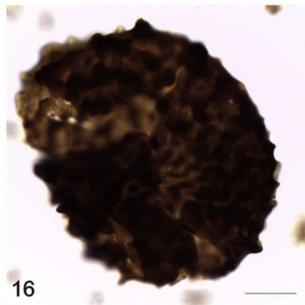
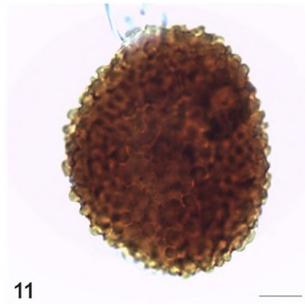
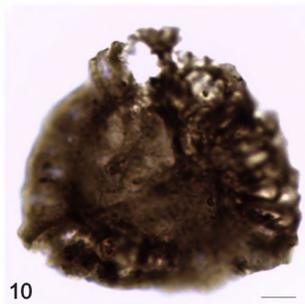
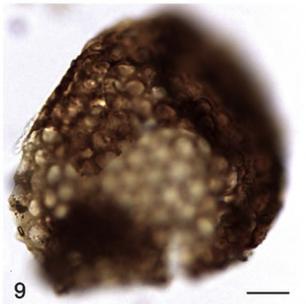
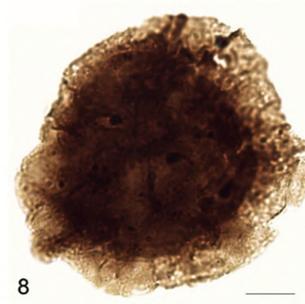
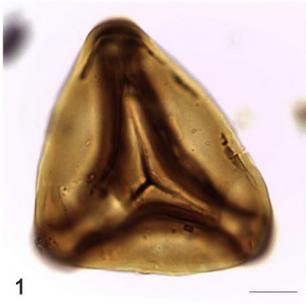


Plate 1

Fig. 5





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Plate 2

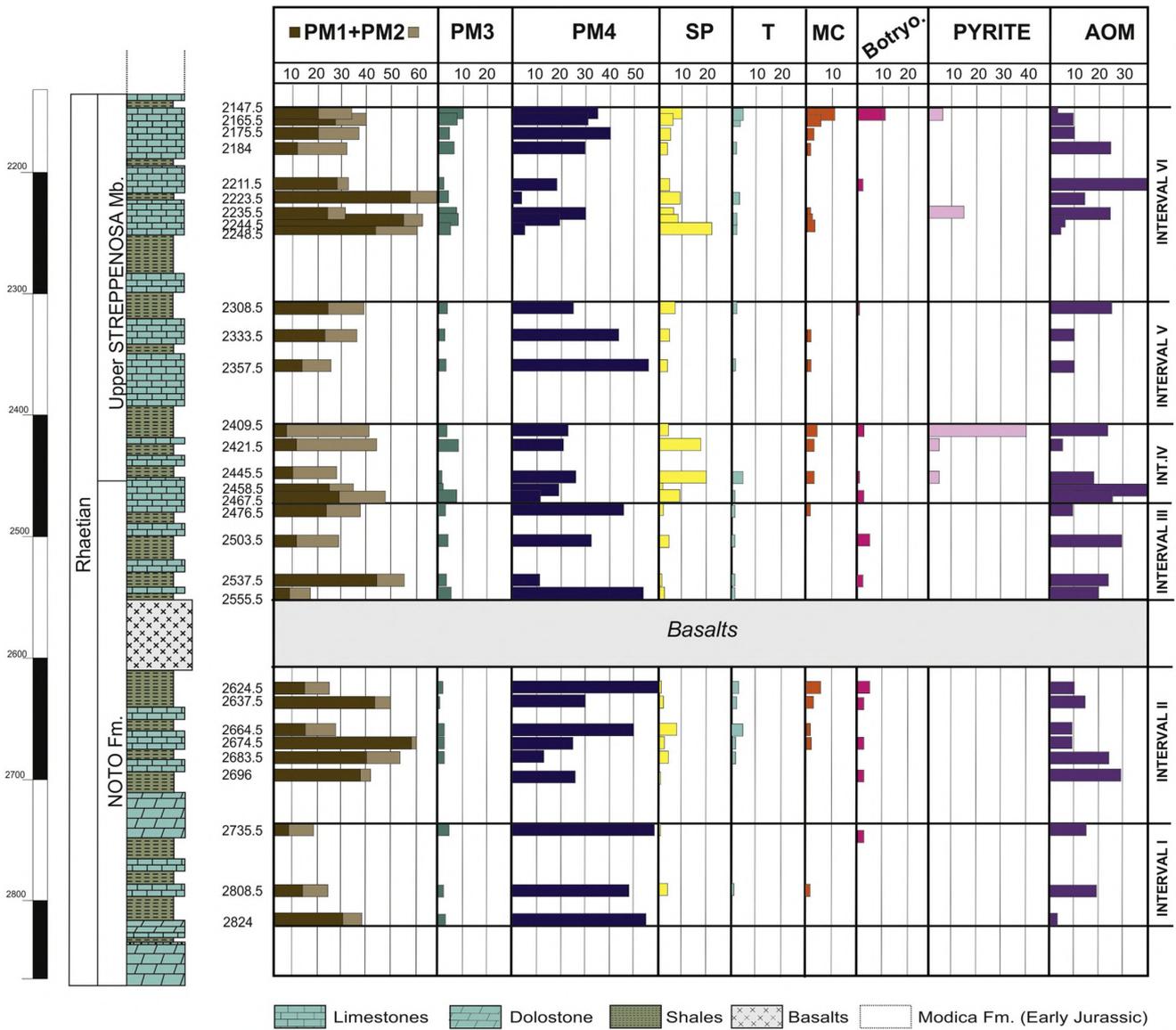


Fig. 6.

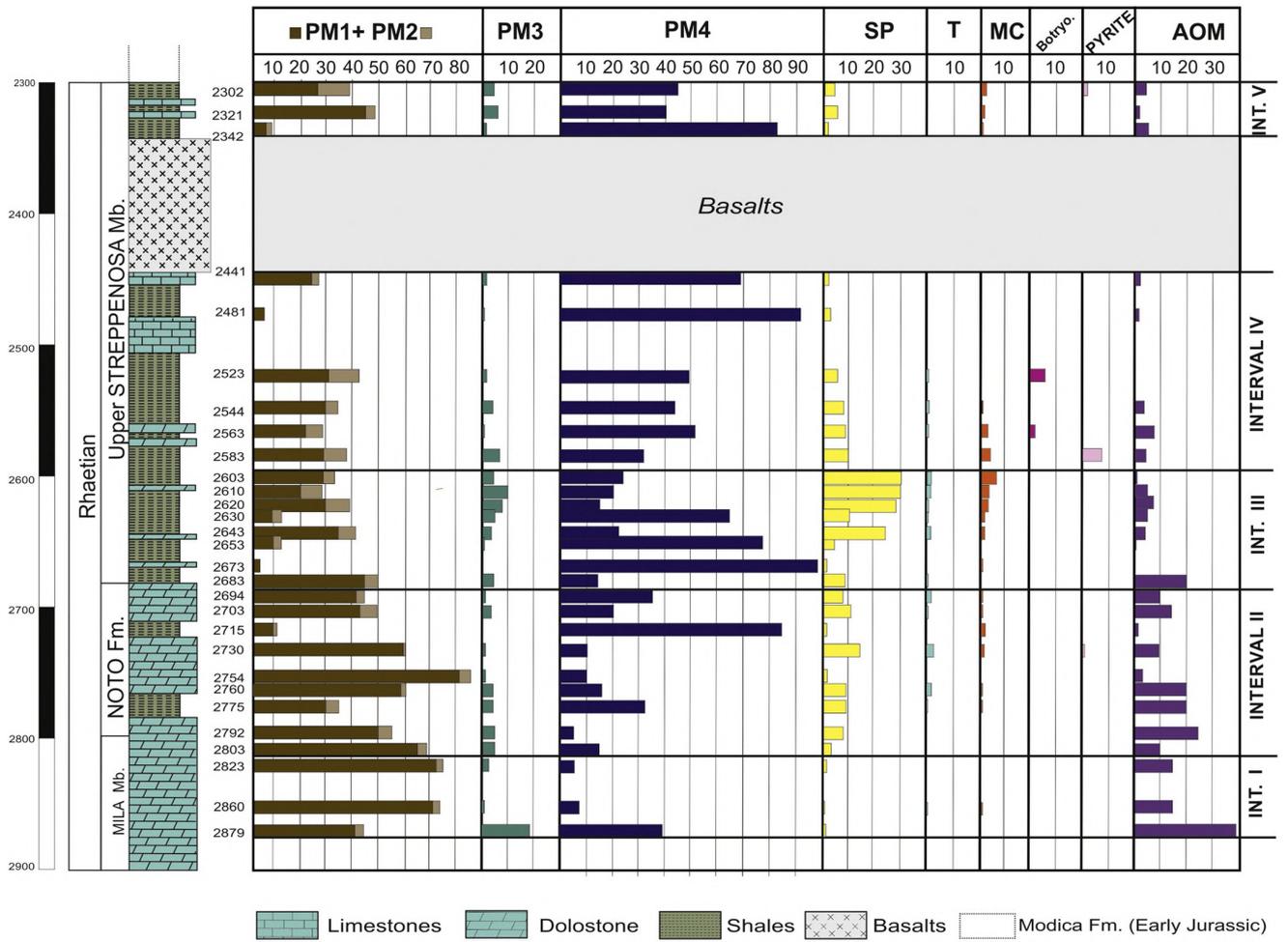


Fig. 7.

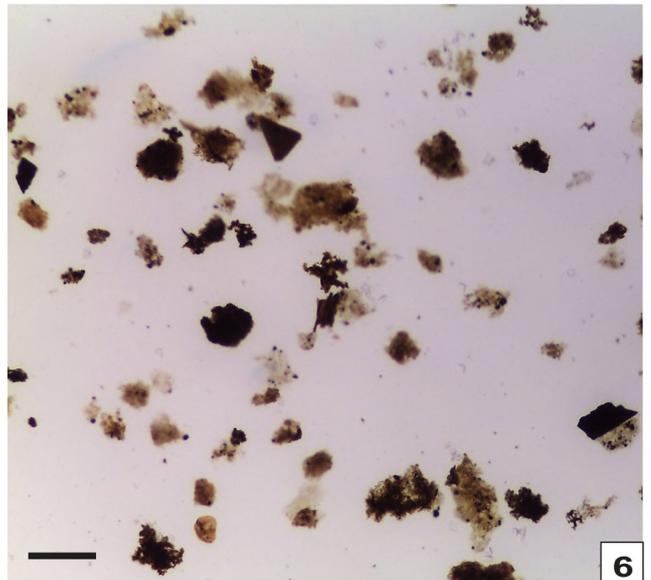
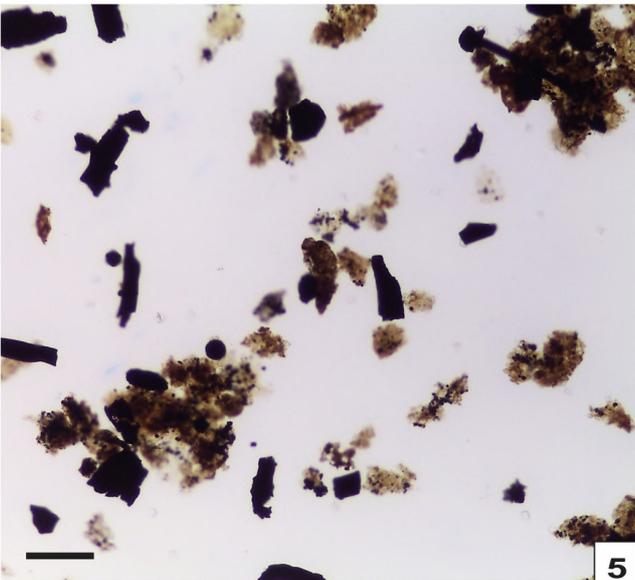
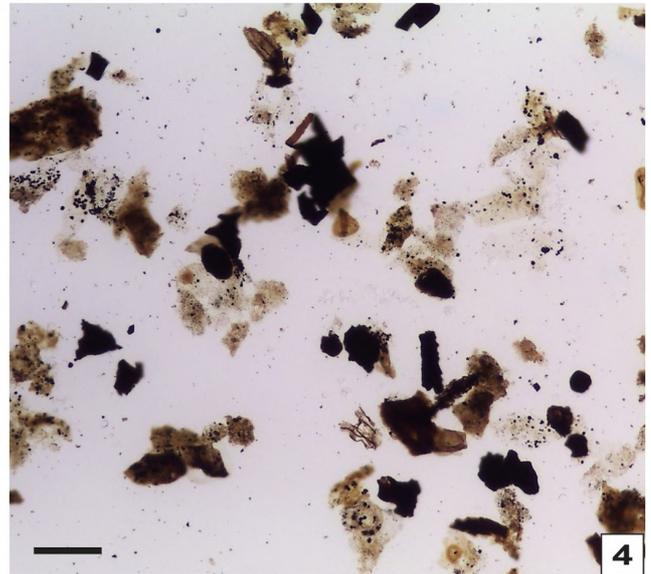
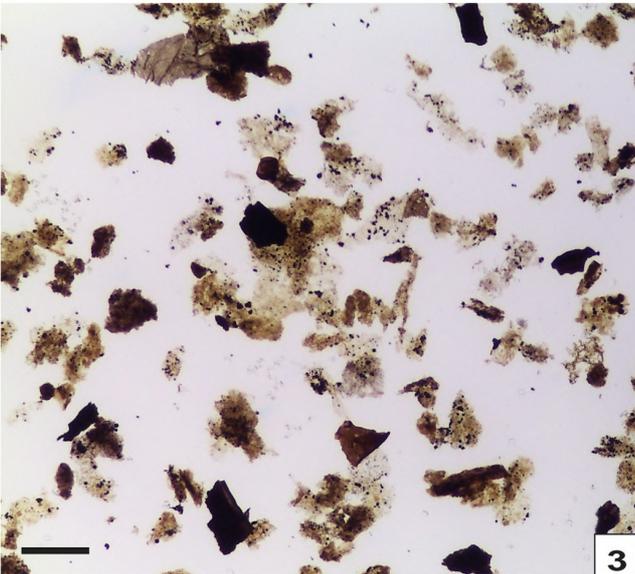
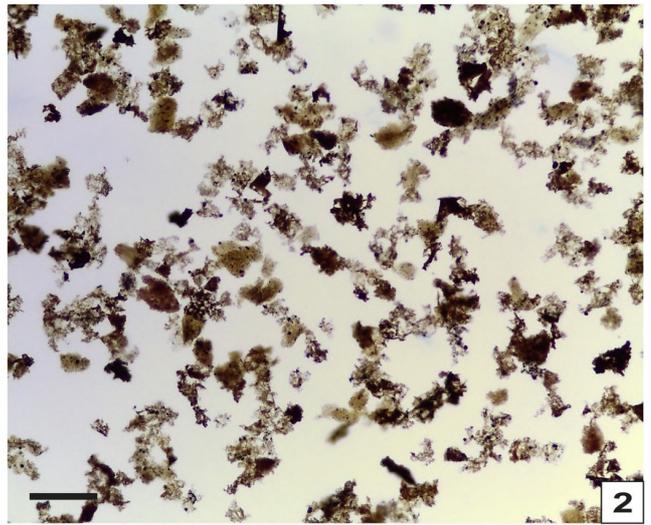
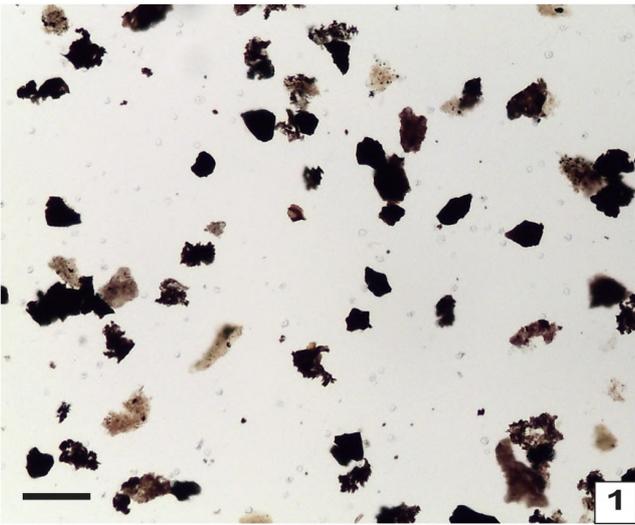


Plate 3

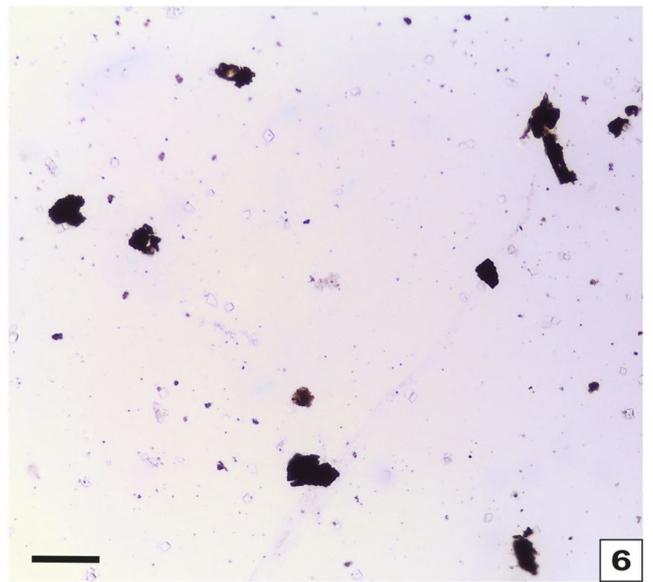
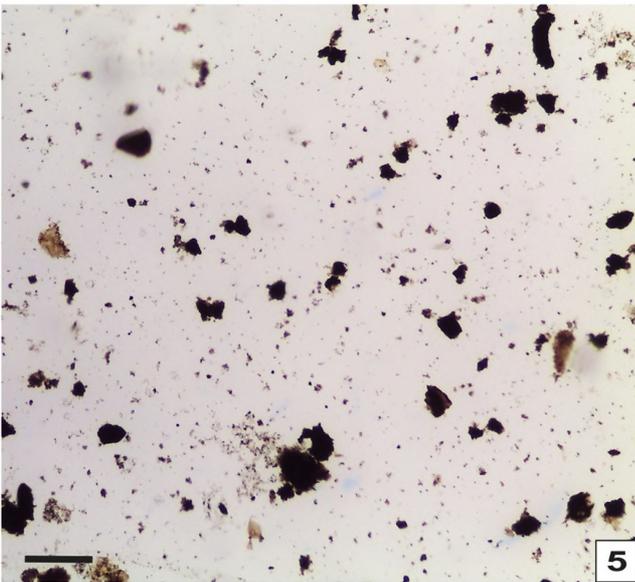
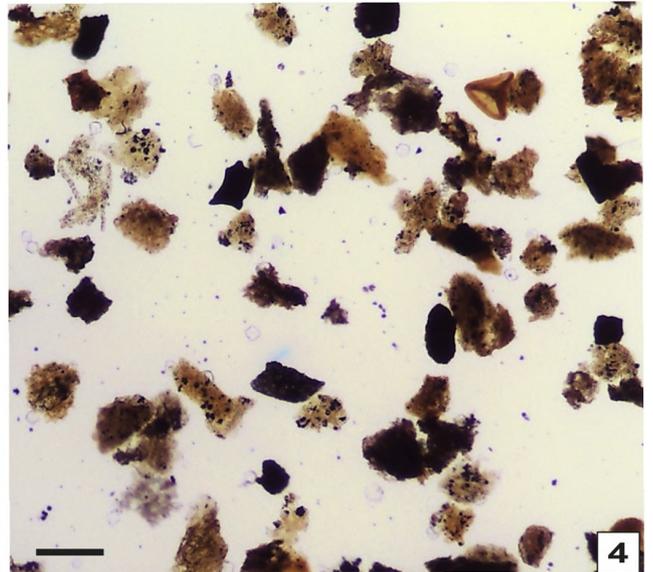
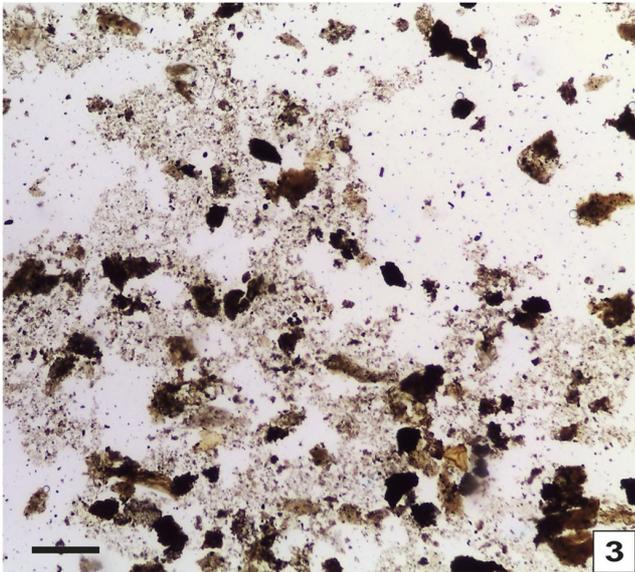
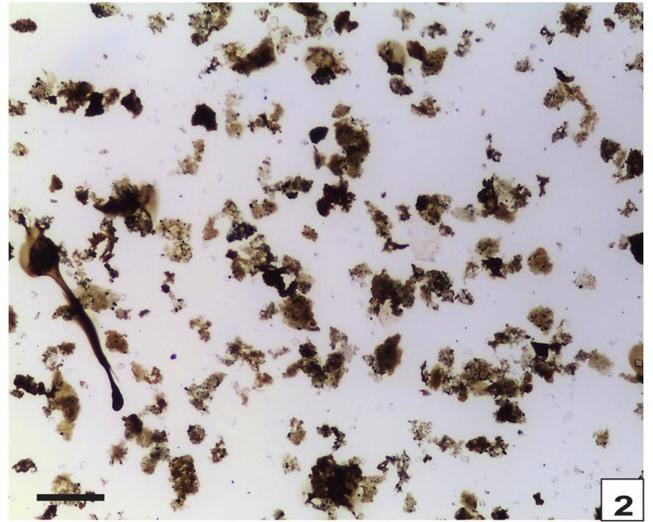
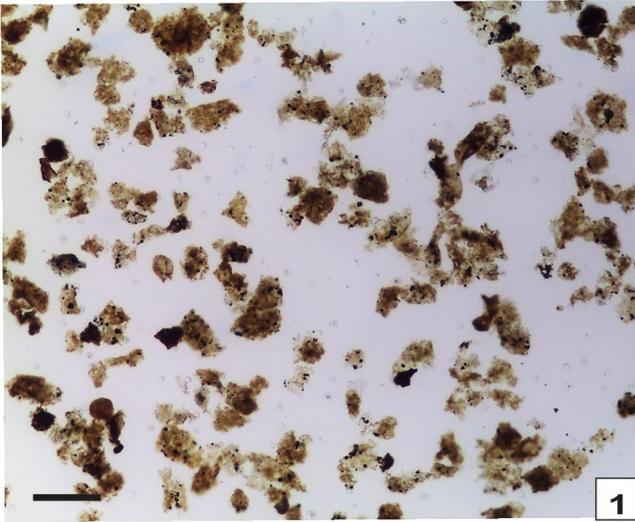


Plate 4