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Palynomorphs from the Devonian Talacasto and Punta Negra Formations,
Argentinean Precordillera: New biostratigraphic approach

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1 **Palynomorphs from the Devonian Talacasto and Punta Negra formations,**
2 **Argentinean Precordillera: new biostratigraphic approach**

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20

21 **Abstract**

22 Marine and terrestrial palynomorphs from the Loma de los Piojos section of the
23 Precordillera Basin, western Argentina, were analyzed. The studied section embraces
24 the upper part of the Lower Devonian Talacasto Formation and the whole Lower-
25 Middle? Devonian Punta Negra Formation, which constitute the Gualilán Group.
26 Terrestrial palynomorphs dominate the Lower Devonian Talacasto Formation whereas
27 marine palynomorphs dominate the Lower-Middle? Devonian Punta Negra Formation.
28 Even though palynomorphs are scarce, poorly preserved, and marked by low diversity,

29 they enable, together with previous palynological data, a new biostratigraphic
30 interpretation for the Devonian units of the Gualilán Group. In addition, an age
31 reassessment of the guide horizon of Keidel (1921), and a correlation of sections from
32 different parts of the basin are provided. Based on palynological data, the age of the
33 Talacasto Formation is constrained to the Lochkovian-late Pragian and the age of the
34 Punta Negra Formation is now considered to be restricted to the late Pragian-Emsian,
35 although a younger age for the uppermost part of Punta Negra is possible.

36

37 **Keywords**

38 Devonian; Argentinean Precordillera; organic-walled phytoplankton; miospores;
39 biostratigraphy.

40

41 **1. Introduction**

42 Paleobiogeographic particularities of the Devonian basins from Southwestern
43 Gondwana have greatly hindered regional or wider biostratigraphic accuracy based on
44 faunas. This region, known as the Malvinokaffric Realm (Richter and Richter, 1942),
45 corresponds to a major marine paleobiogeographic unit developed in circum-polar
46 austral cool/cold water settings, mainly recognizable from the Lower-Middle Devonian
47 of southern South America, South Africa and Antarctica. It is characterized by the
48 virtual absence of typical Paleozoic biostratigraphic guide groups such as conodonts,
49 graptolites and goniatites (Boucot and Racheboeuf, 1993) and the endemism of its
50 faunas in South American basins, except in the northernmost ones. Accordingly,
51 palynostratigraphy has proven to be critical in order to reach biostratigraphic proposals
52 for international comparisons and correlations of the Malvinokaffric sedimentary
53 successions (e.g. Grahn, 2002, 2005; Loboziak and Melo, 2002; Melo and Loboziak,
54 2003; Rubinstein et al., 2005, 2008; Troth, 2011; Grahn et al., 2013). However, in the
55 Precordillera Basin of central-west Argentina, the palynology of the Lower Devonian
56 successions is still poorly known.

57 The Early-Middle Devonian of the Central Argentine Precordillera is represented by
58 the Gualilán Group (Baldis, 1975), which includes the Talacasto (Padula et al., 1967)
59 and the Punta Negra (Bracaccini, 1949) formations. Despite their extensive exposures,
60 the age of these units has been debated for many decades. Hence, the ages of both

61 formations have been proposed based on alternative macrofaunistic groups, mainly
62 brachiopods and trilobites (e.g. Benedetto et al., 1992; Herrera, 1993, 1995; Herrera and
63 Bustos, 2001; Holloway and Rustán, 2012). Even though the information provided by
64 these benthic groups has been relevant, in many cases, comprehensive revisions are
65 needed, particularly taking into account changes in the stratigraphic range of index taxa,
66 derived either from new worldwide findings or from age revisions of the units from
67 which they were originally reported.

68 The first palynological reports from these Devonian units were restricted to the
69 basal part of the Talacasto Formation at Cerro del Fuerte locality (Le Hérisse et al.,
70 1997) and two levels of the Punta Negra Formation in the Talacasto Creek (Rubinstein,
71 1999, 2000). Although these papers represent very relevant biostratigraphic studies in
72 the region, they also include inconsistencies in the species designation that require
73 further revision.

74 Recently, more extensive palynological studies from the Talacasto and Punta
75 Negra formations were carried out at Cerro La Chilca and Talacasto Creek areas (García
76 Muro et al., 2017). The recorded assemblages, dominated by marine organic walled
77 microphytoplankton, allow constraining the age of the Talacasto Formation to the
78 Lochkovian- probably late Pragian and the lower part of the Punta Negra Formation to
79 the early Emsian. Furthermore, the age of the marker horizon identified by Keidel (1921)
80 and Astini (1991) was restricted to the late Pragian-early Emsian.
81 Paleobiogeographically, the record of typical Gondwanan taxa suggested affinities with
82 this paleocontinent.

83 In the present contribution, the upper part of the Talacasto Formation and the
84 complete Punta Negra Formation at Loma de los Piojos, where the thickest Devonian
85 successions crop out, were analyzed. In a reliable stratigraphic framework controlled by
86 the Keidel marker horizon, the record of new palynological data contributes to better
87 constrain the ages and the stratigraphic relationships around the boundary of these units
88 and to accurately date the guide stratigraphic horizon involved.

89 1.1. *Geological setting of the Gualilán Group*

90 The Devonian outcrops of the Gualilán Group, extend mainly along the Central
91 Precordillera, in the Province of San Juan, extend with a north-south alignment for more

92 than 150 km, from Río Jáchal in the north to Río San Juan in the south (Fig. 1). Some
93 additional outcrops, lateral equivalents to the lower part of the Talacasto Formation,
94 were recognized in the Province of La Rioja, nearly 170 km to the north of the Jáchal
95 área (Rubinstein et al., 2010; Rustán and Vaccari, 2010a).

96 The Talacasto Formation overlies the mainly Upper Silurian Los Espejos
97 Formation by a paraconformity (Astini and Maretto, 1996). To the south, the hiatus
98 encompasses a Ludlow to lowest Lochkovian interval. However, to the north, near the
99 Jáchal river the Silurian-Devonian boundary has been recognized in the uppermost
100 interval of the Los Espejos Formation (Benedetto et al., 1992; Carrera et al., 2013;
101 García Muro et al., 2014a; García Muro and Rubinstein, 2015).. The lower part of the
102 unit typically displays Lochkovian to Pragian intensely bioturbated dark mudstone
103 levels (Fig. 2), interpreted as an external muddy platform with no influence of wave
104 action and a low sedimentation rate (Astini, 1991). Towards the top, it passes to sand-
105 rich deposits influenced by wave action (Astini, 1991; Carrera and Rustán, 2015;
106 Rustán and Balseiro, 2016). The Talacasto Formation was considered as a muddy-shelf
107 depositional system, developed during a high stand (Astini, 1991).

108 The upper part of the Talacasto Formation contains a nearly 10 m-thick
109 distinctive ochre, nodule-bearing fossiliferous horizon, interpreted as a marker horizon
110 by Keidel (1921) and Astini (1991). The Punta Negra Formation overlies the Talacasto
111 Formation by a paraconformity that is younger to the north (Salas et al., 2013 and
112 references therein). In the southern sections near the San Juan river, the Punta Negra
113 Formation lies directly above Keidel's horizon through a 40 m-thick interval of
114 alternating purple and green pelites (García Muro et al., 2017). By contrast, in the Loma
115 de los Piojos locality (Fig. 1), the Talacasto Formation extends ca. 300 m above
116 Keidel's horizon (Rustán, 2016), and the overlying Punta Negra Formation includes at
117 the lower part ca. 210 m of light green mudstones. According to the palynological
118 results at the Cerro La Chilca section (García Muro et al., 2017), the layers of the
119 Talacasto Formation located immediately above Keidel's marker horizon would be
120 Pragian to possibly late Pragian in age, in contrast with the previously proposed early
121 Emsian age based on brachiopods (Herrera, 1993, 1995; Rustán and Vaccari, 2010b;
122 Salas et al., 2013; Sterren et al., 2015; Rustán, 2016; Cichowolski and Rustán, 2017).

123 Toward the top, the Punta Negra Formation consists of a coarser and thicker
124 succession of intercalated green to blackish-green sandstones and siltstones (Fig. 2). In
125 the Loma de Los Piojos section, the Punta Negra Formation is not arranged in such
126 typical tabular heterolithic rhythmic layers as in the type section in the San Juan river
127 area. The uppermost part of the unit is covered by the Upper Carboniferous glacial
128 diamictite of the Guandacol Formation (Cuerda, 1965). The depositional environment
129 of the Punta Negra Formation has been extensively discussed and interpreted as
130 deposits of submarine fans (e.g. González Bonorino, 1975), a submarine fan with storm
131 events (Herrera and Bustos, 2001), deltaic deposits (Bustos and Astini, 1997) or shallow
132 low-energy marine deposits evolving towards turbiditic sandstone beds (Edwards et al.,
133 2009).

134 The Loma de los Piojos section is located ca. 8 km southwest of Jáchal city. In
135 this locality, the Talacasto Formation reaches 1145 m, the maximum thickness of the
136 unit. In turn, the Punta Negra Formation is approximately 1000 m thick to the south of
137 the basin whereas in the surveyed transect it is restricted to nearly 350 m due to a strong
138 erosive paleorelief filled with the glacial diamictites representing the base of the
139 Carboniferous Guandacol Formation. On the basis of the trilobite *Acanthopyge*
140 (*Lobopyge*) *balliviani* (Kozłowski, 1923) reported by Rustán and Vaccari (2010a) in the
141 very same transect, the layers located about 250 m above the base of the Punta Negra
142 Formation were considered Middle Devonian in age. Taking into account the lithology,
143 the stratigraphic position and the brachiopod fauna, such as *Salopina* and *Metaplasia*
144 (L. Benedetto, personal communication), these strata were correlated with those
145 reported in a similar stratigraphic position at the southwest Río de las Chacritas section
146 and dated as Emsian by Herrera and Bustos (2001).

147 **2. Materials and Methods**

148 Fifty palynological samples from the Loma de los Piojos locality in an east-west
149 transect situated at 30° 18,504' S/68° 47,399' W were analyzed. Fifteen samples were
150 collected from the upper 250 m of the Talacasto Formation and 35 samples from the
151 lower 320 m of the Punta Negra Formation, that is up to the erosive discordance of the
152 base of the Upper Carboniferous Guandacol Formation (Fig. 2).

153 The samples were processed in the Palaeopalynology Laboratory of IANIGLA,
154 CCT CONICET Mendoza, Argentina, using the standard palynological HCl–HF–HCl

155 acid maceration technique (Traverse, 2007). The organic residue was sieved through a
156 10 µm mesh and, when necessary, oxidized for 1 or 2 minutes with nitric acid (HNO₃).
157 The palynomorphs were mounted in glycerin jelly as permanent palynological slides
158 and examined using light microscopy and differential interference contrast (DIC) when
159 the palynomorphs were transparent.

160 A set of samples from the Punta Negra Formation, which yielded scarce
161 palynomorphs, was also processed at the University of Liège, Belgium. The samples
162 were processed by a similar acid maceration technique, although all of them were
163 oxidized in low-grade Schulze solution (HNO₃ and KClO₃) and sieved through a 12 µm
164 mesh. The samples corresponding to each level are listed in Table 1 and numbered from
165 1 to 50, from the bottom to the top of the section.

166 The slides are housed in the palaeopalynological collection of IANIGLA, CCT
167 CONICET Mendoza, Argentina. Each figured palynomorph is located by slide numbers
168 and England Finder coordinates between brackets.

169 3. Results

170 The Talacasto and the Punta Negra formations at the Loma de los Piojos section,
171 yielded poorly preserved marine and terrestrial palynological assemblages of low
172 diversity. Palynomorph identifications at the species or generic level were possible in a
173 total of 26 levels, and they are illustrated in Figures 3-5. In the other levels, the
174 palynomorphs were carbonized, broken and dark brown to black in color, thus hindering
175 their identification even as a marine or terrestrial group.

176 The studied section of the Talacasto Formation, which ranges from two meters
177 below Keidel's horizon up to the top of the formation, nearly 250 m above such
178 horizon, yielded badly preserved palynomorphs. In most of the levels, only a few
179 palynomorphs, mainly miospores, were identified (Fig. 3). The presence of trilete spores
180 cf. *Verrucosisporites* sp. and ?*Knoxisporites riondae*, in the lowermost studied level is
181 highlighted. The following levels lack biostratigraphically relevant palynomorphs.

182 The uppermost studied level of the Talacasto Formation (level 15), located
183 approximately 230 m above Keidel's horizon, yielded the most abundant and diversified
184 assemblage, with 10 miospore species, four organic-walled microphytoplankton species,
185 and two fresh water acritarchs. Trilete spores that first appear in the Early Devonian, such

186 as *Dibolisporites echinaceus*, *Dictyotriletes emsiensis* Morphon and ?*Knoxisporites*
187 *riondae*, were recognized in this level. This is the only level containing some
188 identifiable marine organic-walled microphytoplankton species, such as *Diexallophasis*
189 *remota* Group and *Veryhachium trispinosum* Group, although they are
190 biostratigraphically widely distributed in Lower Paleozoic rocks and thus not useful to
191 accurately date this part of the section.

192 The Punta Negra Formation yielded a more diverse and better preserved
193 palynomorph assemblage than the Talacasto Formation. The marine organic-walled
194 microphytoplankton was more abundant and diverse than the terrestrial palynomorphs.
195 A few fresh water acritarch taxa are also present (Figs. 4 and 5). The lower light green
196 mudstone-rich interval of approximately 210 m yielded very scarce species such as
197 *Schizocystia saharica* and *Estiastra rugosa*. Upwards in the sequence, in level 32, close
198 to the coarser succession of intercalated greenish-gray wackes and siltstones, some
199 miospores including *Cymbosporites proteus* occur. Levels 33 to 39 contained some
200 biostratigraphically relevant organic-walled microphytoplankton species such as
201 *Bimerga paulae* and *Navifusa bacilla*, and the miospore *Chelinospora verrucata*
202 Morphon. The upper 80 m of the section, below the Carboniferous diamictite of the
203 Guandacol Formation, yielded poorly preserved palynomorphs with no biostratigraphic
204 value.

205 The complete list of species recorded in the section is given in the appendix. The
206 corresponding illustrations are depicted in Figures 6-10.

207 4. Discussion

208 4.1. Biostratigraphy

209 Le Hérisse et al. (1997) studied the lower part of the Talacasto Formation
210 recording more than 50 species with scarce biotratigraphic value. Nevertheless,
211 according to these authors, the chitinozoan *Urochitina lobo* Volkheimer et al., 1986,
212 the acritarchs *Demorhethium lappaceum* Loeblich and Wicander 1974 and *Schizocystia*
213 *pilosa*, and the trilete spores *Dibolisporites echinaceus* and *Dictyotriletes emsiensis*
214 Morphon, would suggest a maximum Lochkovian age for this part of the unit. Later, the
215 Silurian/Devonian boundary was recognized within the underlying Los Espejos
216 Formation by García Muro et al. (2014a) based on palynomorphs. These authors dated
217 the uppermost 6.25 m of the Los Espejos Formation and the basal 31.5 m of the

218 Talacasto Formation at the Río Jáchal locality as Lochkovian. This age is supported by
219 the occurrence of the spores cf. *Streelispora newportensis* (Chaloner and Streel)
220 Richardson and Lister 1969, cf. *Dictyotriletes emsiensis* Morphon, *Cymbosporites*
221 *paulus?* in Wellman (1993) and the phytoplankton species *Schizocystia pilosa* and
222 *Polyedryxium condensum* in the upper part of the Los Espejos Formation. *Streelispora*
223 *newportensis* is one of the index species of the early Lochkovian MN Biozone defined
224 by Richardson and McGregor (1986) for the Old Red Sandstone continent, and also of
225 the equivalent *micrornatus-newportensis* Biozone erected by Streel et al. (1987). This
226 species was also recorded in the Peri-Gondwanan terrane of Spain (Richardson et al.,
227 2001) and the Moesian Terrane (Stemans and Lakova, 2004). The miospores
228 *Dibolisporites echinaceus* and *Breconisporites* sp. from the lower part of the Talacasto
229 section, would be consistent with that age.

230 In a recent contribution of García Muro et al. (2017), a comprehensive revision
231 of previous palynological data was performed, and new palynological assemblages from
232 the Talacasto Formation and the lower part of the Punta Negra Formation, from
233 different localities, were studied. The lower 15 m of the Talacasto Formation at the
234 Cerro La Chilca locality yielded palynomorph taxa suggesting a Lochkovian *s.l.* age,
235 such as *Schizocystia pilosa* and *Winwaloewsia distracta* (Deunff) Deunff 1977, and the
236 trilete spore *Dictyotriletes emsiensis* Morphon. The upper 850 m of the sequence
237 yielded organic-walled microphytoplankton species such as *Palacanthus ledanoisii* and
238 *Polyedryxium* cf. *P. decorum*, and spores such as *Dictyotriletes favosus* McGregor and
239 Camfield 1976 and *?Knoxisporites riondae*, that would indicate a Pragian to late
240 Pragian age. According to these findings, the Lochkovian/Pragian boundary would
241 occur in the lowest 15 m of the Talacasto Formation at the Cerro La Chilca locality.
242 These results would contradict the age assigned to the stratigraphic unit based on fauna
243 evidence, in which the Lochkovian/Pragian boundary was generally located hundreds of
244 meters above the base of the Talacasto Formation and the uppermost part of the unit
245 was assigned to the Emsian (Herrera, 1993, 1995).

246 The lowermost level of the Talacasto Formation studied herein, located 2 m
247 below the guide horizon of Keidel at the Loma de los Piojos locality, yielded a few
248 poorly preserved palynomorphs. However, a specimen of *?Knoxisporites riondae* and a
249 specimen of cf. *Verrucosisporites* sp. were recorded. The oldest record of
250 *?Knoxisporites riondae* comes from the San Pedro Formation, northern Spain, dated as

251 “post-Ludlow-Pridoli?” or, most probably, as earliest Lochkovian by Cramer and Diéz
252 (1975) and Rodriguez (1978). Nevertheless, no megafossil independent age control was
253 available. It is noteworthy that ?*Knoxisporites riondae* is a common species in the
254 Pragian/Emsian boundary in Gondwanan and peri-Gondwanan regions (e.g. Steemans,
255 1989; Breuer and Steemans, 2013).

256 The sample from Keidel’s horizon (level 2) did not provide recognizable
257 palynomorphs, although some phytoclasts and possible miospores were recorded. The
258 remaining section, that consists of the upper 248 m, yielded some miospore specimens,
259 such as cf. *Acinosporites* sp., *Dibolisporites* sp., *Dictyotriletes* sp., and cf.
260 *Dibolisporites echinaceus*. Even though these taxa are already present in the
261 Lochkovian and some of them extend higher in the Devonian, their occurrence would
262 not contradict a possible late Pragian age suggested for the lowermost studied levels.

263 The uppermost studied level (15) from the Talacasto Formation yielded
264 miospores such as *Dictyotriletes emsiensis* Morphon, *Cymbohilates* sp. and
265 *Apiculiretusispora* sp., as well as additional specimens of ?*Knoxisporites riondae* (Fig.
266 3). This assemblage would be consistent with a late Pragian age proposed for the lower
267 levels that were sampled herein, especially due to the presence of ?*Knoxisporites*
268 *riondae* and the lack of taxa that first appear in the Emsian. Moreover, it should be
269 considered that equivalent levels of the Talacasto Formation above Keidel’s horizon at
270 the Cerro La Chilca locality were also interpreted as late Pragian based on
271 palynomorphs (García Muro et al., 2017). Thus, the Emsian age assigned for the upper
272 part of the Talacasto Formation based on faunal evidence (e.g. Herrera, 1993), would
273 not be supported by the palynomorphs recorded at the Loma de los Piojos locality. Le
274 Hérissé et al. (1997) proposed an Emsian age for strata located 112 m above the
275 boundary with the underlying Los Espejos Formation at the northern Cerro del Fuerte
276 locality. However, this age was based on doubtfully classified chitinozoans that need
277 further revision. Consequently, the Emsian age for the Talacasto Formation is not based
278 on trustworthy biostratigraphic data.

279 Rubinstein (1999, 2000) dated the Punta Negra Formation exposed in the
280 northern margin of the Talacasto Creek as Eifelian-Givetian based on badly preserved
281 palynomorphs, of which some were misidentified, such as *Arkonites bilixus* Legault
282 1973, *Emphanisporites annulatus* McGregor 1961 and *Verrucosisporites scurrus*

283 (Taugourdeau-Lantz) Richardson and McGregor 1986 (Rubinstein, 1999, p. 15; figure
284 2). According to Rubinstein (2000), the presence of chitinozoans doubtfully assigned as
285 *Fungochitina pilosa* (Collinson and Scott, 1958) and ?*Ancyrochitina langei* (Sommer
286 and van Boekel, 1964) would point to a middle-late Givetian age. Moreover, the
287 acritarch identified as *Bimerga bensonii* Wood 1995 was considered as an element
288 confirming the Middle Devonian age (Rubinstein, 2000); yet, a taxonomic revision
289 reassigned it to *Bimerga paulae* Le Hérissé 2011, restricting its stratigraphic range to
290 the Pragian-Emsian interval (Le Hérissé, 2011; Daners et al., 2017; Rubinstein et al., in
291 press). Thus, a reappraisal of these assemblages suggests an age not younger than
292 Emsian for the bearing levels. According to García Muro et al. (2017), the lowest part of
293 the Punta Negra Formation (40 m), close to the sample location of Rubinstein (1999,
294 2000), in the Talacasto Creek, would not be older than Early Emsian in age due to the
295 occurrence of marine palynomorphs such as *Navifusa bacilla* (Molyneux et al., 1996;
296 Fatka and Brocke, 2008).

297 The shales and mudstones from the lower part of the Punta Negra Formation at
298 Loma de los Piojos yielded poorly preserved palynomorphs, some of which could be
299 reworked. Such is the case of the cryptospore *Artemopyra brevicosta*, recorded in level
300 17, which ranges worldwide from the Wenlock to the Lochkovian (Steemans et al.,
301 2007). The trilete miospore *Emphanisporites protophanus*, recorded in level 27, could
302 also be reworked, as its younger records correspond to the Pridoli (Steemans et al.,
303 2007). Therefore, the basal part of the formation could not be accurately dated based on
304 the palynomorphs hereby recognized.

305 *Cymbosporites proteus*, recorded in level 32, first appears in the Lochkovian
306 MN Zone (Richardson and McGregor, 1986; Streel et al., 1987; Steemans, 1989) and
307 ranges up to the Emsian (McGregor and Camfield, 1976; Rodriguez, 1978). However,
308 no Emsian stratigraphic markers were found in this part of the unit and, consequently,
309 an age younger than Pragian cannot be assumed.

310 In level 33 from the Punta Negra Formation, two specimens of *Bimerga paulae*
311 were recorded. As stated above, this Gondwanan species would have a short
312 biostratigraphic distribution corresponding to the late Pragian-early Emsian, with
313 records in Brazil (e.g. Le Hérissé, 2011; Mendlowicz Mauller et al., 2007), Bolivia -as
314 *B. bensonii* variant A (Pérez-Leyton, 2007; Wood, 1995)-, Argentina -as *B. bensonii*

315 (Rubinstein, 2000)- and Uruguay (Daners et al., 2017; Rubinstein et al., in press). Le
316 Hérissé (2011) also pointed out that *B. bensonii* would be typically recorded in younger
317 ages, from the late Eifelian to the Givetian strata, and that the specimen of *B. paulae*
318 recognized by Lange (1967) in the Famennian of the Amazonas Basin in Brazil could
319 be considered as reworked. Therefore, the occurrence of *B. paulae* in level 33 would
320 indicate an age restricted to the late Pragian/early Emsian for this part of the unit,
321 approximately 210 m above the base of the Punta Negra Formation in the Loma de los
322 Piojos section (Fig. 2).

323 A single specimen of *Latosporites* sp. 1 in Breuer and Steemans (2013) was
324 recorded upward in the section of the Punta Negra Formation, in level 34. According to
325 these authors, this species first appears in the Eifelian, whereas *L. ovalis* is constrained to
326 the Pragian-Emsian. However, *Latosporites* sp. 1 in Breuer and Steemans (2013) is only
327 known from Saudi Arabia, thus inhibiting its biostratigraphic use. The other recorded
328 species from the same level, such as *Archaeozonotriletes chulus*, *Cymatiosphaera* sp.,
329 *Polyedryxium* sp. and the *Veryhachium trispinosum* Group, are not germane for
330 determining a precise age.

331 *Riculasphaera fissa*, a species formerly considered to be restricted to the
332 Lochkovian (e.g. Loeblich and Drugg, 1968; Loeblich and Wicander, 1976; Wicander,
333 1986; Limachi et al., 1996; Rubinstein et al., 2008; Vavrdová et al., 2011), was
334 recorded in level 35. This species was also recognized in the Pragian of the Talacasto
335 Formation (García Muro et al., 2017); thus, this findings would confirm a younger
336 stratigraphic range for this species. Interestingly, Turnau and Racki (1999) illustrate a
337 *Pterospermella* sp. (Plate IV, 15; p. 260) in the Givetian of Poland that actually seems
338 to correspond to a *Riculasphaera fissa* specimen, similar to those illustrated by
339 Rubinstein et al. (2008; fig. 7, 4; p. 180).

340 Few biostratigraphically relevant species, such as *Navifusa bacilla* in level 37,
341 *?Knoxisporites riondae* in level 38 and *Chelinospora verrucata* Morphon in level 39,
342 appear upward in the section. *Navifusa bacilla* is known in Devonian strata since the
343 Emsian (Molyneux et al., 1996; Fatka and Brocke, 2008; Grahn et al., 2013) and *C.*
344 *verrucata* Morphon last appears in the Emsian (McGregor and Camfield, 1976; García-
345 Muro et al., 2014b and references therein). Thus, the stratigraphic interval from levels
346 37 to 39 of the Punta Negra Formation at the Loma de los Piojos locality, approximately

347 at 230 m above the base of the Punta Negra Formation, would be restricted to the
348 Emsian.

349 Level 40 corresponds to the location in which the trilobite *Acanthopyge*
350 *bolliviani* was collected (Rustán and Vaccari, 2010a; Salas et al., 2013). This taxon is
351 considered Middle Devonian in South America (Salas et al., 2013 and references
352 therein). Even though the few and poorly preserved spores herein recognized, such as
353 cf. *Acinosporites* sp., *Dibolisporites* sp. and *Dictyotriletes* sp., are not useful to
354 accurately date this level, the occurrence of *Schizocystia pilosa* would not support an
355 age younger than Early Devonian (García Muro et al., 2017 and references therein).

356 The upper part of the section contains palynomorphs that would point to an
357 Early Devonian age. The acritarch species *Veryhachium trispininflatum*, and the
358 miospore *Raistrickia* sp., are the most relevant taxa from level 41. The uppermost levels
359 yielded a few badly preserved palynomorphs with no biostratigraphic value, such as
360 *Lophosphaeridium* sp. and *Veryhachium trispinosum* Group.

361 4.2. Keidel's guide horizon and age correlations of the Devonian stratigraphic units

362 The new palynological information provided here has significant implications,
363 particularly regarding the age and the stratigraphic relationships of Keidel's guide
364 horizon and the boundary between the Talacasto and the Punta Negra formations.

365 Keidel's guide horizon was interpreted to be located around the Pragian-Emsian
366 boundary. This horizon, which constitutes the top of the Talacasto Formation at its type
367 locality at the Talacasto Creek, was initially referred to the earliest Emsian by Herrera
368 (1993, 1995). Later, a late Pragian age for this bed, otherwise indicative of the top of
369 assemblage biozone B of Herrera, was favored by Herrera (1995) based on the
370 brachiopod *Australostrophia penoensis* Herrera, 1995. However, an early Emsian age
371 was not excluded when considering the evolutionary stage of development of the
372 median septum in this taxon. The presence of the chitinozoan *Bulbochitina bulbosa*
373 (Paris 1981) in the upper part of the Talacasto Formation, in the Talacasto Creek, was
374 mentioned by Le Hérisse et al. (1997). This finding would support a late Pragian age for
375 this part of the unit because *Bulbochitina bulbosa* is the index species of the
376 homonymous Interval Range Biozone (Paris et al., 2000). The palynological evidences
377 provided in this contribution also excludes a possible Emsian age for Keidel's horizon,

378 supporting therefore a late Pragian age for such stratigraphic reference, very useful in
379 the field. Additionally, this bed was correlated (Rustán and Vaccari, 2010b) with the
380 seismic “marker” 4 of the SG VII from the base of the Emsian in Bolivia (Albariño et
381 al., 2002, p. 53), which has been related to a sudden rise of relative sea level after a
382 strong eustatic fall displayed during the Pragian (Albariño et al., 2002; Álvarez et al.,
383 2003).

384 This age constraining of the Keidel’s bed, enables alternative interpretations of
385 the stratigraphic relationships between units conforming the Gualilán Group. According
386 to the most accepted stratigraphic scheme, as shown in recent faunalistic contributions
387 by Rustán and collaborators, layers immediately overlying the Keidel’s bed are laterally
388 equivalents despite the changes of facies. However, recent palynological evidence
389 suggests different ages for the layers overlying the Keidel’s bed from south to north. An
390 Emsian age was suggested for the purple and green pelites in the southern Talacasto
391 Creek area corresponding to the base of the Punta Negra Formation. Although, at
392 northern sections, a probably late Pragian age is assumed for the uppermost greenish
393 mudstones in the Cerro La Chilca section (García Muro et al, 2017) and for the greenish
394 gray siltstones and very fine sandstones at the Loma de los Piojos section (this
395 contribution), both corresponding to the upper part of the Talacasto Formation. In this
396 last interpretation, a stratigraphic hiatus immediately above the Keidel’s bed should be
397 invoked to explain the diachronism in layers in such a stratigraphic position along the
398 basin.

399 This alternative hypothesis suggests that the base of the Punta Negra Formation
400 would not be so broadly diachronic, so that the lowest purple and green pelites from the
401 Talacasto Creek would be lateral equivalents to the green pelitic lowest interval of the
402 Punta Negra Formation at the Loma de los Piojos section (levels 16 to 32 in this
403 contribution). Meanwhile, they would not have equivalent stratigraphic records in the
404 Cerro La Chilca section due to the erosion by the basal discordance of the
405 Carboniferous Guandacol Formation. This could explain the absence of late Pragian
406 layers overlying Keidel’s horizon in the Talacasto Creek locality, but its occurrence in
407 the Cerro La Chilca section (García Muro et al., 2017), and to the north at the Loma de
408 los Piojos section (Fig. 11). Further work is necessary in order to confirm this new
409 stratigraphic proposal. Although based on scarce palynological information, it will have
410 important implications for the classic brachiopod biostratigraphic scheme of Herrera

411 (1993, 1995). The pioneering biostratigraphic proposals by Herrera allowed the
412 recognition of Lochkovian and Pragian ages in the Precordillera strata of the Gualilán
413 Group that was previously considered exclusively Emsian. New palynological
414 observations suggest that, particularly the Pragian, would be extensively represented.

415 Additional implications of data presented herein involve the Middle Devonian
416 age proposed for the Punta Negra Formation in the Loma de los Piojos section, based on
417 the occurrence of the trilobite *Acanthopyge (Lobopyge) balliviani* nearly 245 m above
418 the base of the unit (Rustán and Vaccari, 2010a; Salas et al., 2013). Although the few
419 and badly preserved palynomorphs herein recorded would not provide enough
420 evidences to support a Middle Devonian age for this part of the section. In contrast, a
421 possible Emsian age should not be excluded. Such a putative Emsian assignment would
422 imply the first Lower Devonian reference for the trilobite *A. (L.) balliviani*, otherwise
423 considered a Middle Devonian guide fossil in Bolivia. Besides, it would confirm the
424 Emsian age based on brachiopods by Herrera and Bustos (2001) from equivalent layers
425 in the Río de las Chacritas section, to the southwest of the basin.

426 5. Conclusions

427 The palynomorphs from the upper part of the Talacasto Formation, above
428 Keidel' horizon, and the complete Punta Negra Formation, both from the Loma de los
429 Piojos locality, were studied.

430 The upper part of the Talacasto Formation mainly yielded terrestrial
431 palynomorphs that would suggest an age not younger than late Pragian.

432 The Punta Negra Formation contained a more diverse palynological assemblage,
433 with more marine than terrestrial taxa. The lower pelitic part of the Punta Negra
434 Formation yielded a few and badly preserved palynomorphs consistent with the late
435 Pragian age proposed for the underling Talacasto Formation. Upwards in the section, in
436 level 33, the late Pragian/early Emsian *Bimerga paulae* appears. The record of *Navifusa*
437 *bacilla*, *?Knoxisporites riondae*, and *Chelinospora verrucata* Morphon in the interval
438 between levels 37 and 39 would confirm an Emsian age for this part of the unit. The
439 palynological assemblages of the uppermost part of the stratigraphic section did not
440 provide evidence for the age of this part of the unit.

441 Accordingly, based on palynological evidence, the age of the Keidel guide
442 horizon is now constrained to the late Pragian. In a newly proposed stratigraphic
443 interpretation, the stratigraphic interval immediately above Keidel's horizon, would be
444 Emsian in the southern sections, near Talacasto Creek, corresponding to the base of
445 Punta Negra Formation, and to the late Pragian in the northern sections, near Jáchal,
446 corresponding to the upper part of the Talacasto Formation.

447 Although the new biostratigraphic palynological data presented herein,
448 encourage new stratigraphic interpretations for the units conforming the Gualilán
449 Group, biostratigraphic information is still scarce and preliminary. Further work is still
450 necessary to definitively date and correlate these formations in different parts of the
451 Precordillera basin.

452

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- 821
- 822 Appendix
- 823 Organic-walled microphytoplankton
- 824 *Ammonidium* sp. (Fig. 7.B)
- 825 cf. *Baculatireticulatus* sp. (Fig. 7.G)
- 826 *Bimerga paulae* Le Hérisse 2011 (Fig. 7.I and 7.J)
- 827 *Cordobesia orientalis* Pöthé de Baldis 1977 (Fig. 7.L)
- 828 *Cymatiosphaera* aff. *C. ledburica* in Mullins (2001) (Fig. 7.M)
- 829 *Cymatiosphaera mirabilis* Deunff 1959 (Fig. 7.N)
- 830 *Cymatiosphaera multisepta* Deunff 1955 (Fig. 7.O)
- 831 *Cymatiosphaera prismatica* (Deunff) Deunff 1961 (Fig. 8.O)
- 832 *Cymatiosphaera salopensis* Mullins 2001 (Fig. 7.P)
- 833 *Cymatiosphaera* spp.
- 834 *Dictyotidium dictyotum* (Eisenack) Eisenack 1955
- 835 *Dictyotidium* cf. *D. variatum* Playford 1977 (Fig. 8.A)

- 836 *Dictyotidium* sp. (Fig. 8.B)
- 837 *Diexallophasis remota* Group Mullins 2001
- 838 *Divietipellis* sp. (Fig. 8.C and 8.D)
- 839 *Dorsennidium rhomboidium* (Downie) Stancliffe and Sarjeant 1996 (Fig. 8.E)
- 840 *Dorsennidium* spp.
- 841 *Dupliciradiatum tenue* González, Moreno and Playford 2005 (Fig. 8.F)
- 842 *Duvernaysphaera aranaides* Cramer 1964 (Fig. 8.G)
- 843 cf. *Duvernaysphaera wilsonii* Deunff 1964 (Fig. 8.H)
- 844 *Estiastra culcita* Wicander 1974 (Fig. 8.K)
- 845 *Estiastra* sp. in Ottone (1996) (Fig. 8.L)
- 846 *Estiastra* sp. (Fig. 8.M)
- 847 *Evittia sanpetrensis* (Cramer) Lister 1970 (Fig. 8.N)
- 848 *Fimbriaglomerella divisa* Loeblich and Drugg 1968 (Fig. 8.P)
- 849 *Gorgonisphaeridium* sp. (Fig. 8.Q)
- 850 *Hoegklintia* cf. *H. longispina* Pöthé de Baldis 1998 (Fig. 8.S)
- 851 *Leiofusa berneseae* Cramer 1964 (Fig. 9.B)
- 852 *Leiosphaeridia* spp.
- 853 *Lophosphaeridium* spp.
- 854 *Melikeriopalla polygonia* (Staplin) Mullins 2001
- 855 *Micrhystridium* spp. (Fig. 6.L)
- 856 *Multiplicisphaeridium* cf. *M. rochesterense* (Cramer and Díez) Eisenack et al. 1973
857 (Fig. 9.C)
- 858 *Multiplicisphaeridium* sp. (Fig. 6.M)

- 859 *Navifusa bacilla* (Deunff) Playford 1977 (Fig. 9.E)
- 860 *Neoverhachium carminae* (Cramer) Cramer 1970
- 861 *Palacanthus ledanoisii* (Deunff) Playford 1977 (Fig. 9.F)
- 862 *Palacanthus stelligerum* (Deunff) Stancliffe and Sarjeant 1996 (Fig. 9.G)
- 863 ?*Polyedryxium asperum* Cramer 1964 (Fig. 9.H)
- 864 *Polyedryxium embudum* Cramer 1964 (Fig. 9.K)
- 865 *Polyedryxium* cf. *P. condensum* Deunff 1971 (Fig. 9.I)
- 866 *Polyedryxium* cf. *P. decorum* Deunff 1955 (Fig. 9.J)
- 867 *Polyedryxium evolutum* Deunff 1955 (Fig. 9.L)
- 868 *Polyedryxium helenaster* Cramer 1964
- 869 *Polyedryxium robustum* Deunff 1971 (Fig. 9.M)
- 870 *Polyedryxium simplex* Deunff 1955 (Fig. 9.N)
- 871 *Polyedryxium* sp. (Fig. 9.O)
- 872 *Pterospermella hermosita* (Cramer) Fensome et al. 1990 (Fig. 9.P)
- 873 *Pterospermella* spp.
- 874 *Quadraditum fantasticum* Cramer 1964 (Fig. 9.Q)
- 875 cf. *Riculasphaera fissa* Loeblich and Drugg 1968 (Fig. 10.B)
- 876 *Schismatosphaeridium longhopense* Dorning 1981 (Fig. 10.C)
- 877 *Schismatosphaeridium* sp. B in Le Hérisse (1989) (Fig. 10.D)
- 878 *Stellinium micropolygonale* (Stockmans and Williere) Playford 1977 (Fig. 10.H)
- 879 *Stellinium rabians* (Cramer) Eisenack et al. 1976 (Fig. 10.I)
- 880 *Verhachium lairdii* Group sensu Servais et al. 2007
- 881 *Verhachium trispiniflatum* Cramer 1964 (Fig. 10.N)

- 882 *Veryhachium trispinosum* Group Servais et al. 2007 (Fig. 6.P)
- 883 *Villosacapsula setosapellicula* (Loeblich 1970) Loeblich and Tappan 1976 (Fig. 10.O)
- 884 cf. *Visbysphaera* sp. A in Richards and Mullins (2003) (Fig. 10.P)
- 885 Fresh water acritarchs
- 886 *Quadrisporites horridus* (Hennelly) Potonié and Lele 1961 (Fig. 9.R)
- 887 *Quadrisporites variabilis* (Cramer) Ottone and Rossello 1996 (Fig. 9.S)
- 888 *Quadrisporites* sp. (Fig. 10.M)
- 889 *Schizocystia pilosa* Jardiné et al. 1972 (Fig. 10.E)
- 890 *Schizocystia saharica* Jardiné et al. 1974 (Fig. 10.F)
- 891 Miospores
- 892 *Acinosporites* sp. (Fig. 7.A)
- 893 cf. *Acinosporites* sp. (Fig. 6.A)
- 894 *Ambitisporites avitus* Morphon Steemans et al. (1996) (Fig. 7.C)
- 895 *Ambitisporites tripapillatus* Moreau-Benoit 1976 (Fig. 7.D)
- 896 *Ambitisporites* sp. (Fig. 6.B)
- 897 *Apiculiretusispora plicata* (Allen) Streeel 1967 (Fig. 7.E)
- 898 *Apiculiretusispora* sp. (Fig. 6.C)
- 899 *Archaeozonotriletes chulus* (Cramer) Richardson and Lister 1969
- 900 cf. *Archaeozonotriletes chulus* (Cramer) Richardson and Lister 1969 (Fig. 6.D)
- 901 *Archaeozonotriletes* spp.
- 902 *Artemopyra brevicosta* Burgess and Richardson 1991 (Fig. 7.F)
- 903 *Brochotriletes foveolatus* Naumova 1953 (Fig. 7.H)
- 904 *Chelinospora verrucata* Morphon García Muro et al. (2014b) (Fig. 7.K)

- 905 *Chelinospora* spp.
- 906 *Cymbohilates* sp. (Fig. 6.E)
- 907 *Cymbosporites proteus* McGregor and Camfield 1976 (Fig. 7.Q)
- 908 cf. *Cymbosporites verrucosus* Richardson and Lister 1969 (Fig. 7.R)
- 909 *Cymbosporites* spp. (Fig. 6.F; Fig. 7.S)
- 910 *Dibolisporites echinaceus* (Eisenack) Richardson 1965 (Fig. 6.G)
- 911 *Dibolisporites* spp. (Fig. 6.H)
- 912 *Dictyotriletes emsiensis* Morphon Rubinstein et al. (2005) (Fig. 6.I)
- 913 *Dictyotriletes* sp. (Fig. 6.J)
- 914 *Emphanisporites protophanus* Richardson and Ioannides 1973 (Fig. 8.I)
- 915 *Emphanisporites rotatus* (McGregor) McGregor 1973 (Fig. 8.J)
- 916 *Gneudnaspora divellomedia* (Chibrikova) Balme 1988 var. minor Breuer et al. 2007
- 917 *Hispanaediscus* cf. *H. verrucatus* (Cramer) Burgess and Richardson 1991 (Fig. 8.R)
- 918 ?*Knoxisporites riondae* Cramer and Díez 1975 (Fig. 6.K and 8.T)
- 919 *Latosporites* sp. 1 in Breuer (2007) (Fig. 9.A)
- 920 *Raistrickia* sp. (Fig. 9.T)
- 921 *Retusotriletes* spp. (Fig. 10.A)
- 922 cf. *Retusotriletes* sp. (Fig. 6.N)
- 923 cf. *Scylaspora* sp. (Fig. 10.G)
- 924 *Synorisporites lybicus* Richardson and Ioannides 1973 (Fig. 10.J)
- 925 *Synorisporites verrucatus* Richardson and Lister 1969 (Fig. 10.K)
- 926 *Tetrahedraletes medinensis* var. *parvus* (Strother and Traverse) Burgess 1991 (Fig.
927 10.L)

928 cf. *Verrucosiporites* sp. (Fig. 6.O)

929

930 Captions

931 **Table. 1.** Samples studied per level. Identification number for the samples prepared in
932 the University of Liège, Belgium and in the Palaeopalynology Laboratory of IANIGLA,
933 CCT CONICET Mendoza, Argentina. Fm.: Formation.

934 **Fig. 1.** Location and geological map of the study area of Loma de los Piojos. Map
935 modified from García Muro and Rubinstein (2015) and García Muro et al. (2017).

936 **Fig. 2.** Stratigraphic section of the upper part of the Talacasto Formation and the Punta
937 Negra Formation at the Loma de los Piojos locality, with the locations of studied
938 samples.

939 **Fig. 3.** Stratigraphic distribution of organic-walled microphytoplankton, fresh water
940 acritarchs (f.w.a.) and miospores in the Talacasto Formation at the Loma de los Piojos
941 section. The black rectangles indicate species presence; the grey rectangles are
942 questionable occurrences.

943 **Fig. 4.** Stratigraphic distribution of the organic-walled microphytoplankton in the Punta
944 Negra Formation at the Loma de los Piojos section. The black rectangles indicate
945 species presence; the grey rectangles are questionable occurrences. *: in Le Hérisse
946 (1989); -: in Ottone (1996); +: in Richards and Mullins (2003); °: in Mullins (2001).

947 **Fig. 5.** Stratigraphic distribution of fresh water acritarchs (f.w.a.) and miospores in the
948 Punta Negra Formation at the Loma de los Piojos section. The black rectangles indicate
949 species presence; the grey rectangle are questionable occurrences. ●: in Breuer (2007).

950 **Fig. 6.** Recorded species from the Talacasto Formation.

951 (A) cf. *Acinosporites* sp., sample 3-9932D (W25/3)

952 (B) *Ambitisporites* sp., sample 15-9943E (G46/2)

953 (C) *Apiculiretusispora* sp., sample 15-9943E (M27/3)

954 (D) cf. *Archaeozonotriletes chulus*, sample 12-9941D (Z39)

- 955 (E) *Cymbohilates* sp., sample 15-9943E (K25/1)
- 956 (F) *Cymbosporites* sp., sample 15-9943E (G30)
- 957 (G) *Dibolisporites echinceus*, sample 15-9943E (O43/3)
- 958 (H) *Dibolisporites* sp., sample 15-9943C (B32)
- 959 (I) *Dictyotriletes emsiensis* Morphon, sample 15-9943D (G34/2)
- 960 (J) *Dictyotriletes* sp., sample 15-9943E (L38/4)
- 961 (K) ?*Knoxisporites riondae*, sample 15-9943E (M27)
- 962 (L) *Micrhystridium* sp., sample 15-9943E (F28/3)
- 963 (M) *Multiplicisphaeridium* sp., sample 15-9943E (F28/3)
- 964 (N) cf. *Retusotriletes* sp., sample 15-9943D (L32/4)
- 965 (O) cf. *Verrucosisporites* sp., sample 1-9930D (O26/2)
- 966 (P) *Veryhachium trispinosum* Group, sample 15-9943E (K34/2)
- 967 Scale bar = 20 μ m.
- 968 **Fig. 7.** Recorded species from Punta Negra Formation
- 969 (A) *Acinosporites* sp., sample 41-73812 (C33/3)
- 970 (B) *Ammonidium* sp., sample 37-73824 (Q50)
- 971 (C) *Ambitisporites avitus* Morphon, sample 36-73823 (K31/4)
- 972 (D) *Ambitisporites tripapillatus*, sample 37-73824 (S38/4)
- 973 (E) *Apiculiretusispora plicata*, sample 37-73829 (J32/1)
- 974 (F) *Artemopyra brevicosta*, sample 17-73782 (X34/2)
- 975 (G) cf. *Baculatireticulatus* sp., sample 34-73788 (O45/3)
- 976 (H) *Brochotriletes foveolatus*, sample 41-73819 (E34/1)
- 977 (I, J) *Bimerga paulae*, sample 33-73893 (E34/4), 33-73822 (Q36/2)

- 978 (K) *Chelinospora verrucata* Morphon, sample 39-73826 (K29/1)
- 979 (L) *Cordobesia orientalis*, sample 33-10091C (J21/2)
- 980 (M) *Cymatiosphaera* aff. *C. ledburica*, sample 36-73823 (O48/4)
- 981 (N) *Cymatiosphaera mirabilis*, sample 36-73823 (J50)
- 982 (O) *Cymatiosphaera multisepta*, sample 37-73829 (S48/2)
- 983 (P) *Cymatiosphaera salopensis*, sample 33-73822 (G41/2)
- 984 (Q) *Cymbosporites proteus*, sample 32-73787 (C35/1)
- 985 (R) cf. *Cymbosporites verrucosus*, sample 38-73825 (T37)
- 986 (S) *Cymbosporites* sp., sample 36-73823 (E36/1)
- 987 Scale bar = 20 μ m.
- 988 **Fig. 8.** Recorded species from Punta Negra Formation
- 989 (A) *Dictyotidium variatum*, sample 37-73824 (D39/3)
- 990 (B) *Dictyotidium* sp., sample 32-73787 (S40/4)
- 991 (C) *Divietipellis* sp., sample 33-73894 (H25/4)
- 992 (D) *Divietipellis* sp., sample 37-73824 (J34)
- 993 (E) *Dorsennidium rhomboidium*, sample 32-73787 (U43/1)
- 994 (F) *Dupliciradiatum tenue*, sample 33-73894 (R26/4)
- 995 (G) *Duvernaysphaera aranaides*, sample 32-73787 (J33/4)
- 996 (H) cf. *Duvernaysphaera wilsonii*, sample 37-73824 (C34/2)
- 997 (I) *Emphanisporites protophanus*, sample 27-73833 (M32/1)
- 998 (J) *Emphanisporites rotatus*, sample 41-73812 (F35/1)
- 999 (K) *Estiastra culcita*, sample 33-73894 (U29)
- 1000 (L) *Estiastra* sp. in Ottone (1996), sample 33-10091C (V43/1)

- 1001 (M) *Estiastra* sp., sample 36-73823 (D37/1)
- 1002 (N) *Evittia sanpetrensis*, sample 37-73824 (U49/2)
- 1003 (O) *Cymatiosphaera prismatica*, sample 37-10095E (W22)
- 1004 (P) *Fimbriaglomerella divisa*, sample 32-73786 (U38/1)
- 1005 (Q) *Gorgonisphaeridium* sp., sample 36-73823 (X36/2)
- 1006 (R) *Hispanaediscus* cf. *H. verrucatus*, sample 36-73823 (S47/1)
- 1007 (S) *Hoegkintia* cf. *H. longispina*, sample 33-73894 (W48/3)
- 1008 (T) ?*Knoxisporites riondae*, sample 40-73814 (O31/1)
- 1009 Scale bar = 20 μ m.
- 1010 **Fig. 9.** Recorded species from Punta Negra Formation
- 1011 (A) *Latosporites* sp. 1, sample 34-73788 (T40)
- 1012 (B) *Leiofusa bernesgae*, sample 37-73824 (F41/2)
- 1013 (C) *Multiplicisphaeridium* cf. *M. rochesterense*, sample 37-73829 (K30/1)
- 1014 (D) *Multiplicisphaeridium* sp., sample 36-73823 (V33)
- 1015 (E) *Navifusa bacilla*, sample 37-73824 (U43/4)
- 1016 (F) *Palacanthus ledanoisii*, sample 37-73824 (S45/4)
- 1017 (G) *Palacanthus stelligerum*, sample 41-73812 (X46/1)
- 1018 (H) ?*Polyedryxium asperum*, sample 37-73824 (N48/3)
- 1019 (I) *Polyedryxium* cf. *P. condensum*, sample 35-73790 (H29/4)
- 1020 (J) *Polyedryxium* cf. *P. decorum*, sample 37-73824 (P44/3)
- 1021 (K) *Polyedrydium embudum*, sample 36-73823 (F45/4)
- 1022 (L) *Polyedryxium evolutum*, sample 36-73823 (M48)
- 1023 (M) *Polyedryxium robustum*, sample 36-73823 (P36)

- 1024 (N) *Polyedryxium simplex*, sample 36-73823 (S28/2)
- 1025 (O) *Polyedryxium* sp., sample 36-73828 (P39/3)
- 1026 (P) *Pterospermella hermosita*, sample 33-73894 (N36/1)
- 1027 (Q) *Quadraditum fantasticum*, sample 27-73833 (X31/2)
- 1028 (R) *Quadrisporites* cf. *Q. horridus*, sample 35-73790 (R49/1)
- 1029 (S) *Quadrisporites variabilis*, sample 37-73824 (J33/1)
- 1030 (T) *Raistrickia* sp., sample 41-73812 (K41)
- 1031 Scale bar = 20 μ m.
- 1032 **Fig. 10.** Recorded species from Punta Negra Formation
- 1033 (A) *Retusotriletes* sp., sample 41-73812 (E48)
- 1034 (B) cf. *Riculasphaera fissa*, sample 35-73790 (Q51/2)
- 1035 (C) *Schismatosphaeridium longhopense*, sample 27-73833 (X31/3)
- 1036 (D) *Schismatosphaeridium* sp. B, sample 33-73827 (U35/1)
- 1037 (E) *Schizocystia pilosa*, sample 40-73814 (R34/1)
- 1038 (F) *Schizocystia saharica*, sample 35-73790 (V38/2)
- 1039 (G) cf. *Scylaspora* sp., sample 33-73894 (M45/4)
- 1040 (H) *Stellinium micropolygonale*, sample 32-73786 (K35)
- 1041 (I) *Stellinium rabians*, sample 37-73824 (M44/3)
- 1042 (J) *Synorisporites lybicus*, sample 27-73833 (O51)
- 1043 (K) *Synorisporites verrucatus*, sample 27-73833 (M37/3)
- 1044 (L) *Tetraedraletes medinensis* var. *parvus*, sample 27-73833 (O30/3)
- 1045 (M) *Quadrisporites* sp., sample 35-73789 (R38/1)
- 1046 (N) *Veryhachium trispininflatum*, sample 41-73812 (O41/3)

1047 (O) *Villosacapsula setosapellicula*, sample 37-73824 (G34)

1048 (P) cf. *Visbysphaera* sp. A, sample 33-73827 (G29/2)

1049 Scale bar = 20 μm .

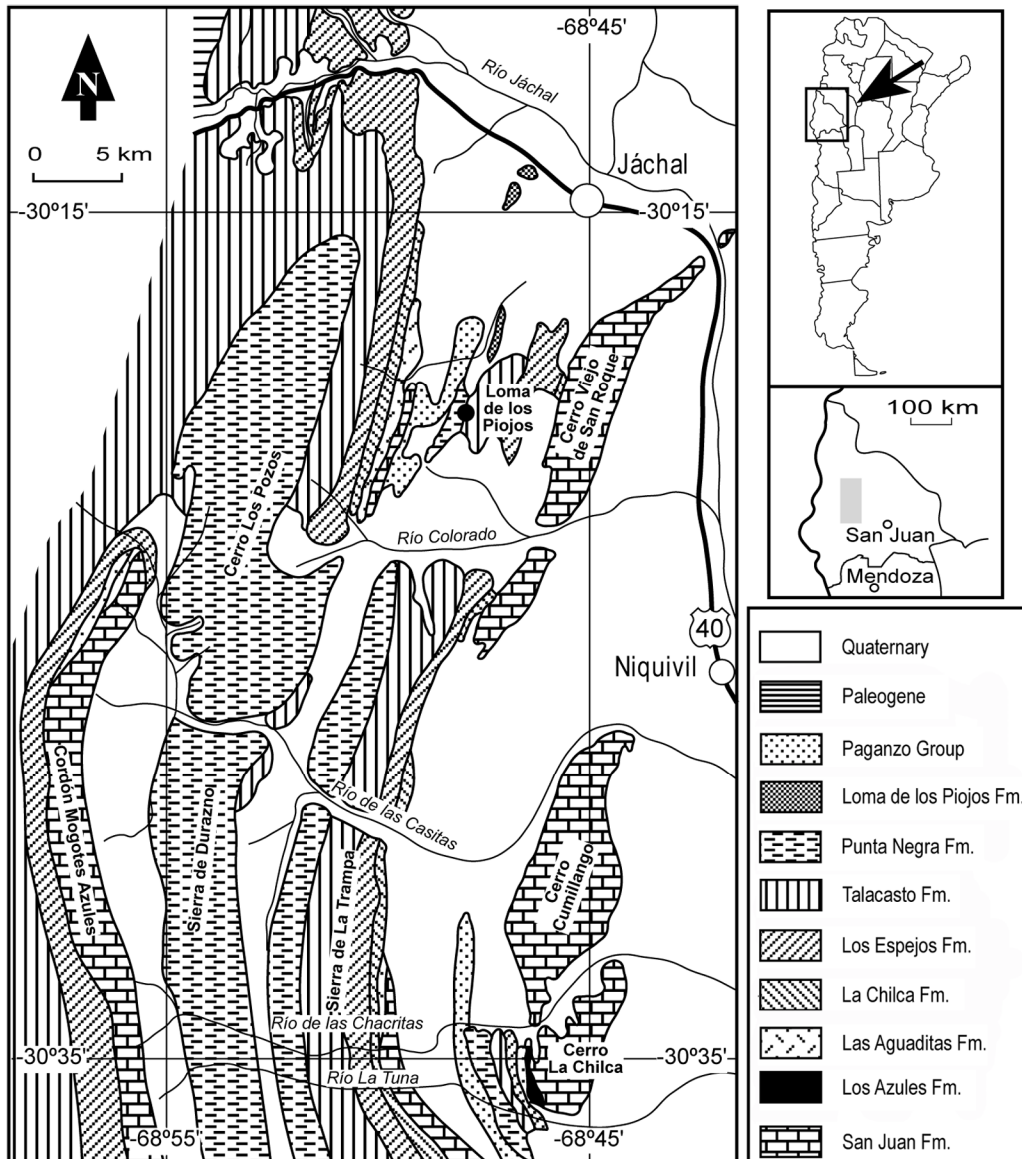
1050 **Fig. 11.** Correlation scheme of the studied sections of the Gualilán Group based on
1051 phytoplankton and miospores assemblages (García Muro et al., 2017 and this
1052 contribution). Loch.: Lochkovian; E: Emsian.

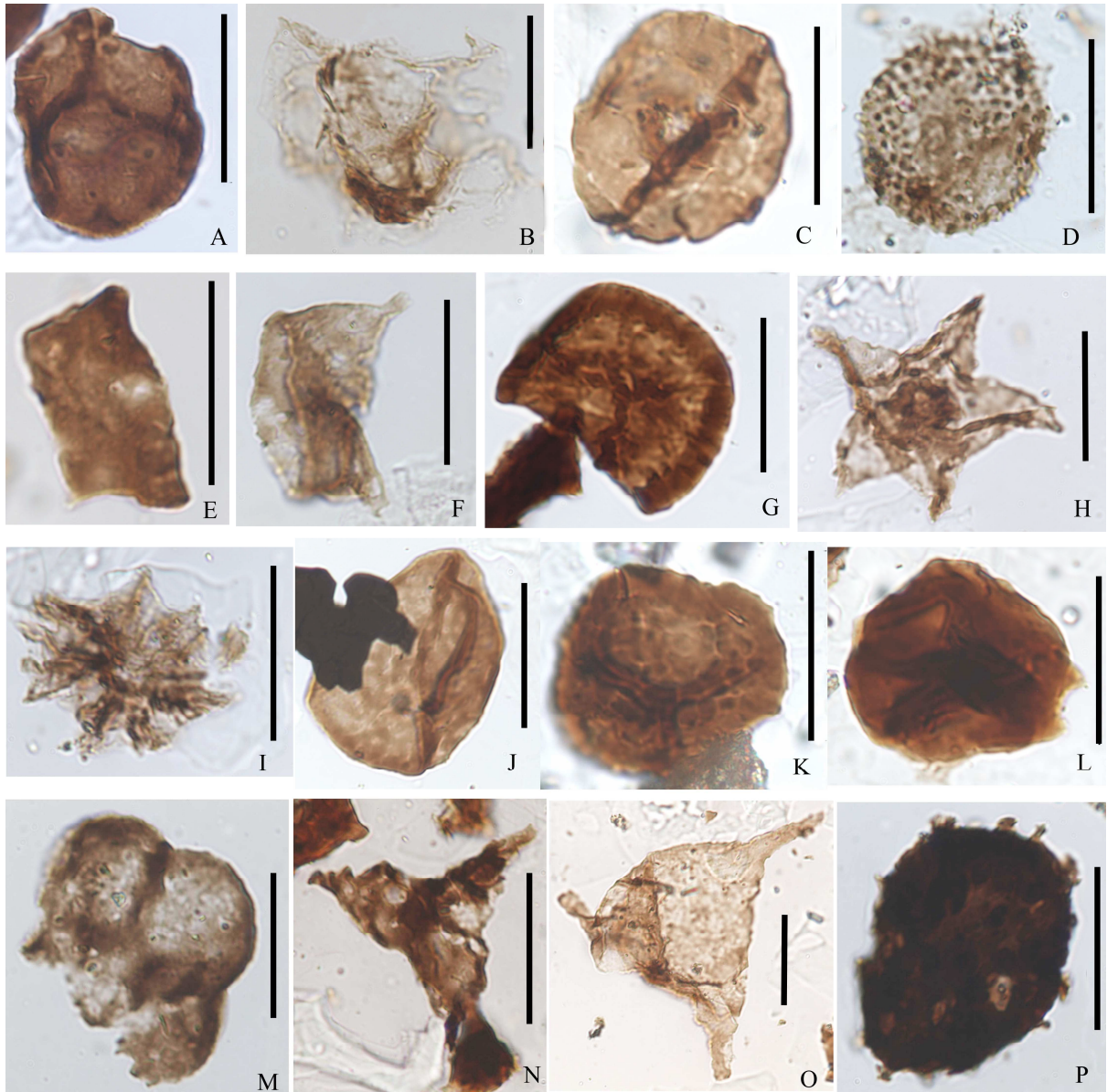
Fm.	Level here simplified	Argentina	Belgium
Talacasto	1	9930	
	2	9931	
	3	9932	
	4	9933	
	5	9934	
	6	9935	
	7	9936	
	8	9937	
	9	9938	
	10	9939	
	11	9940	
	12	9941	
	13	9942	
	14	9960	
	15	9943	
Punta Negra	16	10076	
	17	10074	73782 73783
	18	10075	
	19	10077	
	20	10078	73815
	21	10079	
	22	10080	73816
	23	10081	
	24	10082	73817
	25	10083	73818

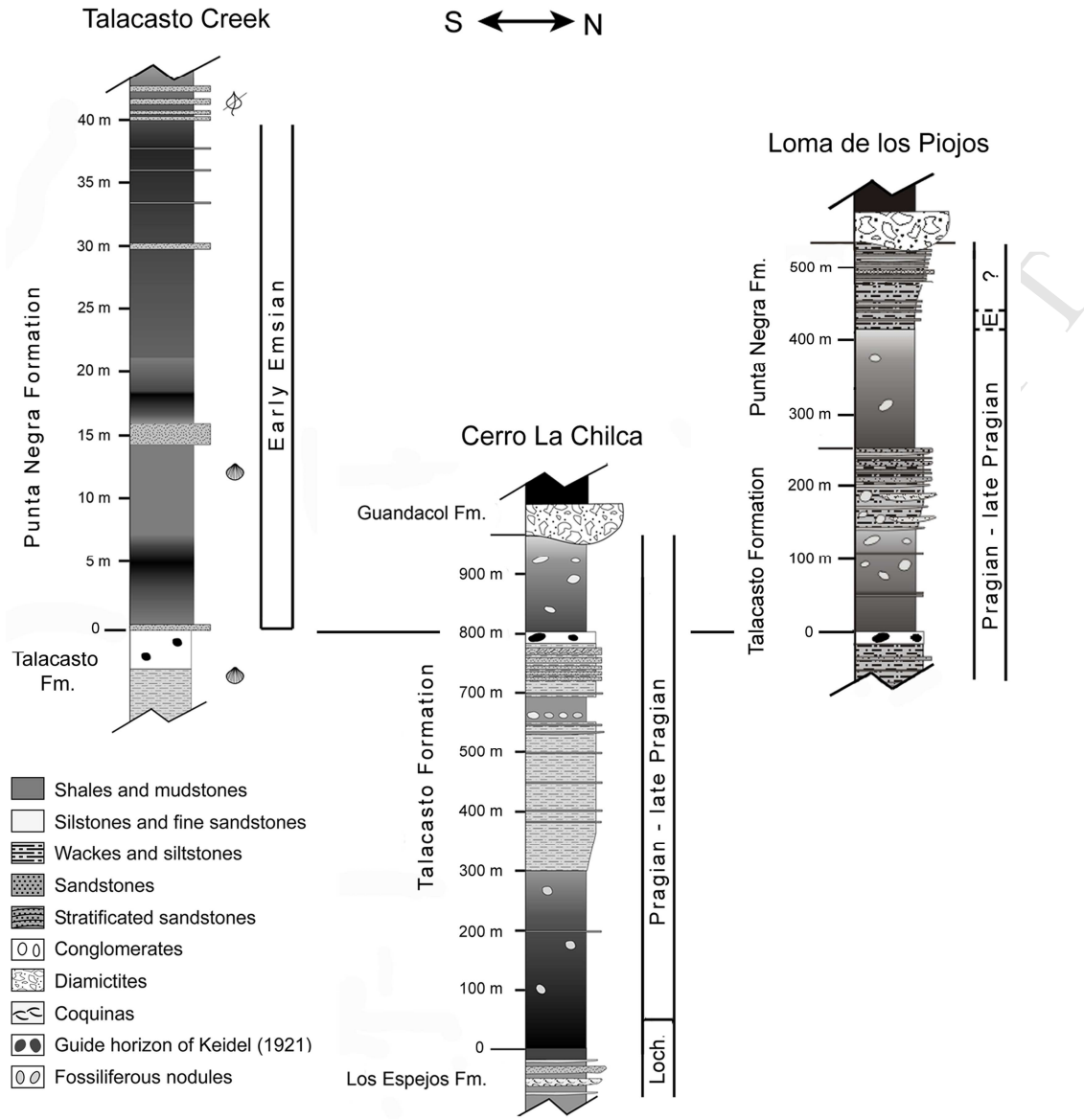
26	10084	
27	10085	73833
28	10086	73784
29	10087	
30	10088	73821
31	10089	73785
32	10090	73786 73787 73791 73792
33	10091	73822 73827 73894 73893
34	10092	73788
35	10093	73789 73790
36	10094	73823 73828
37	10095	73824 73829
38	10096	73825 73830
39	10097	73826
40	10108	73814
41	10098	73812 73813 73819 73820
42	10099	73831
43	10100	73832
44	10101	
45	10102	73834
46	10103	
47	10104	
48	10105	73835
49	10106	

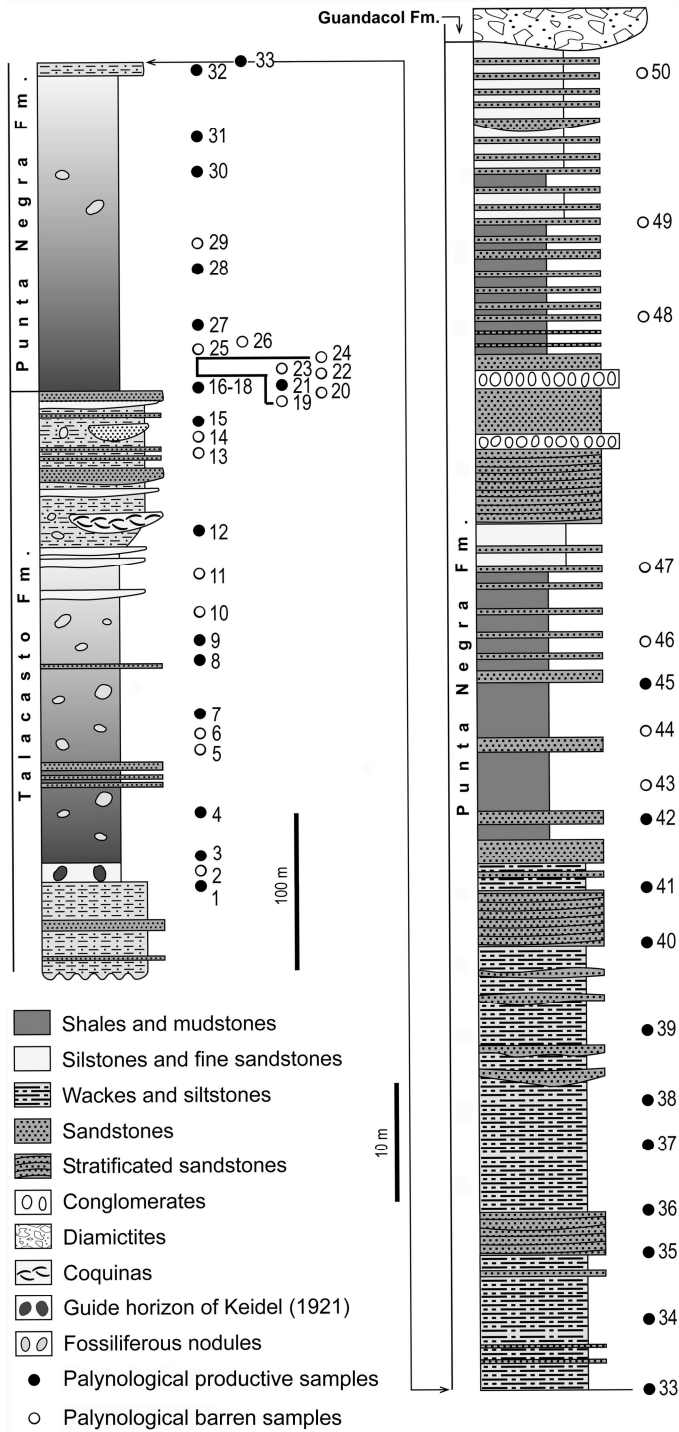
	50	10107	73836
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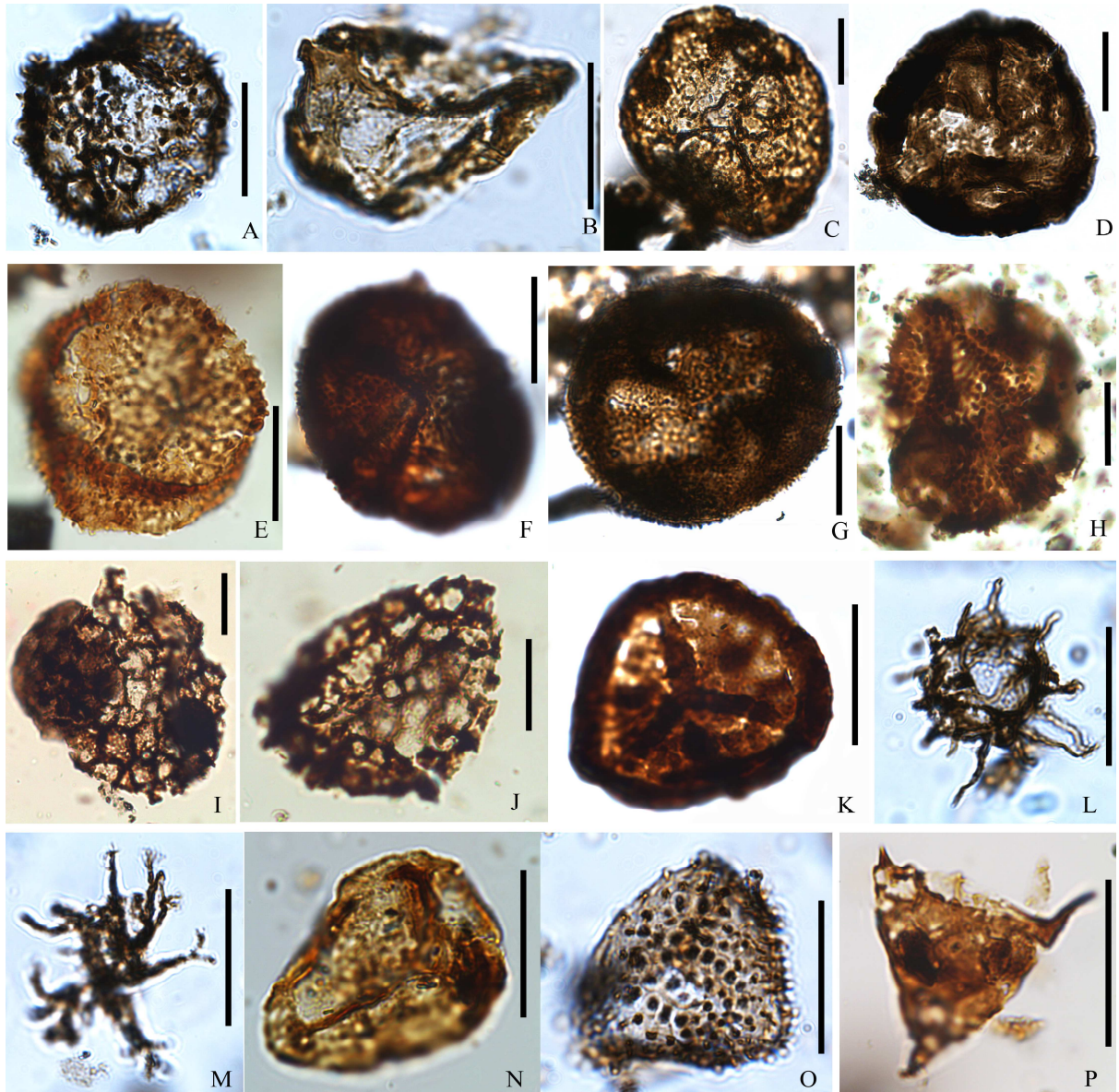


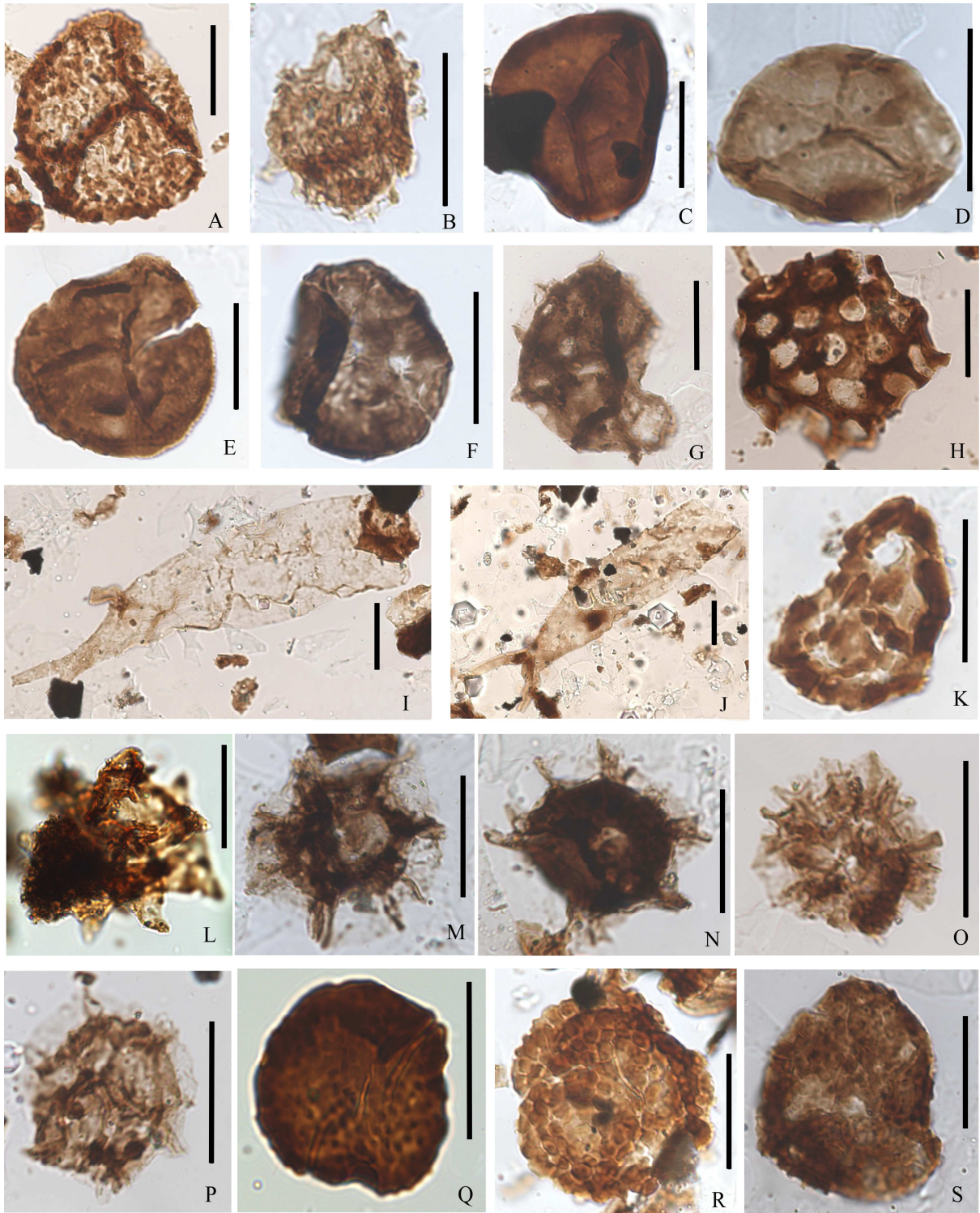


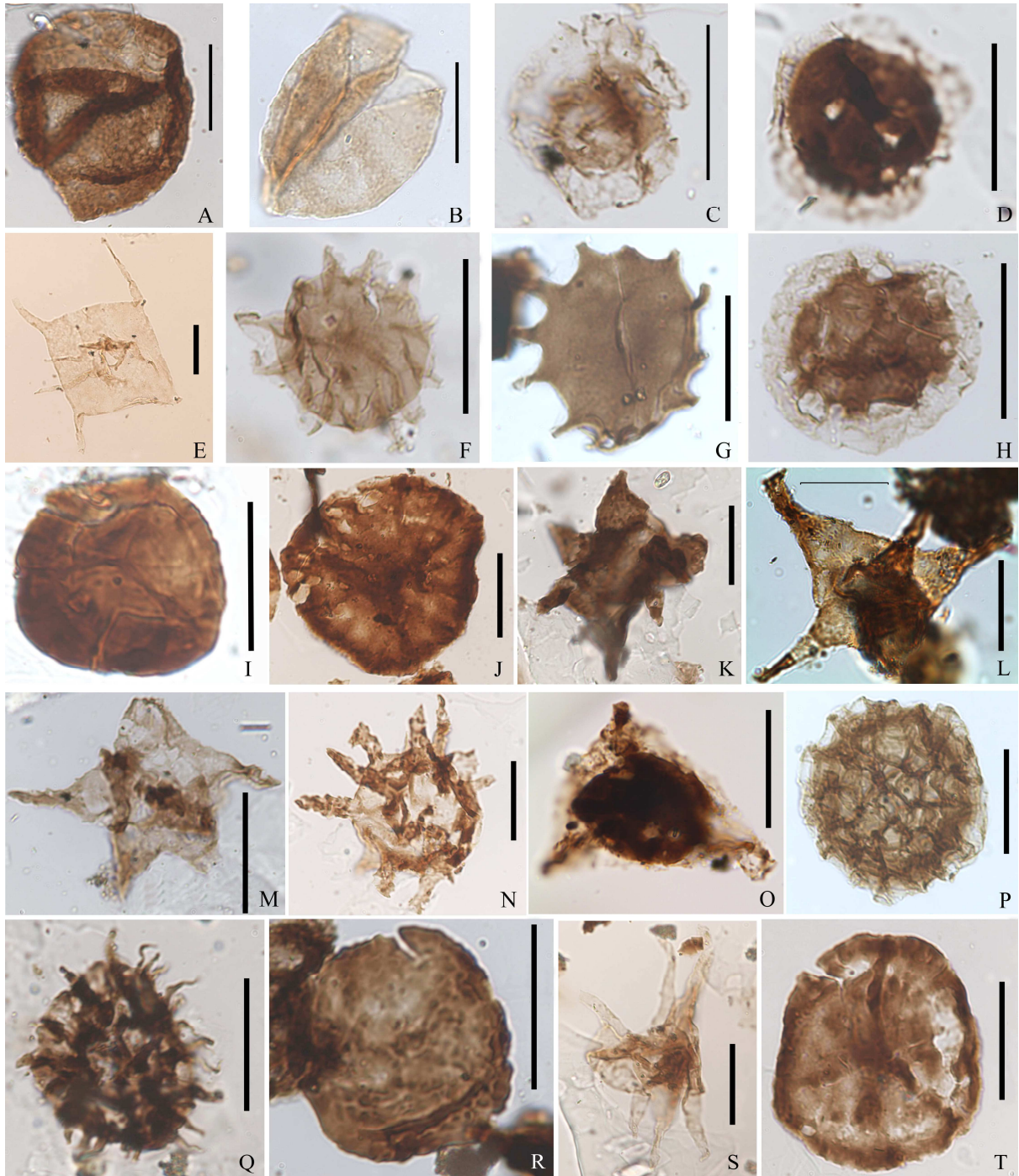


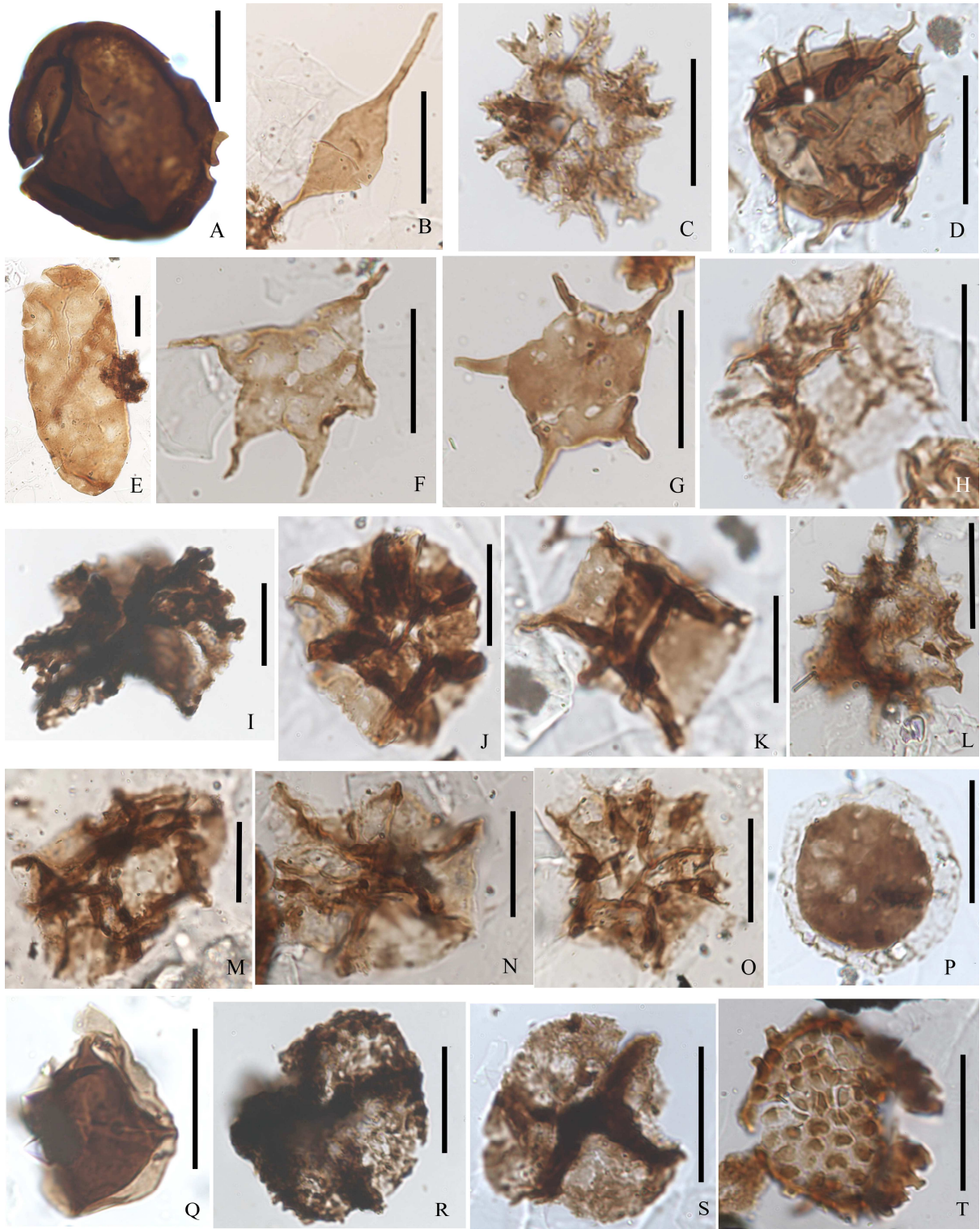
Species	Level	Punta Negra Formation																	
		17	21	27	28	30	31	32	33	34	35	36	37	38	39	40	41	42	45
<i>Veryhachium trispinosum</i> Group																			
<i>Lophosphaeridium</i> spp.																			
<i>Cymatiosphaera</i> spp.																			
<i>Diexallophasis remota</i> Group																			
<i>Pterospermella</i> spp.																			
<i>Leiosphaeridia</i> spp.																			
<i>Micrhystridium</i> spp.																			
<i>Quadratum fantasticum</i>																			
<i>Schismatosphaeridium longhopense</i>																			
<i>Melikeriopalla polygonia</i>																			
<i>Dictyotidium</i> spp.																			
<i>Polyedryxium</i> spp.																			
<i>Dictyotidium</i> cf. <i>D. variatum</i>																			
<i>Dorsennidium</i> spp.																			
<i>Dorsennidium rhomboidium</i>																			
<i>Duvernaysphaera aranaides</i>																			
<i>Fimbriaglomerella divisa</i>																			
<i>Stellinium micropolygonale</i>																			
<i>Veryhachium lairdii</i> Group																			
<i>Gorgonisphaeridium</i> spp.																			
<i>Divietipellis</i> spp.																			
<i>Evittia sanpetrensis</i>																			
<i>Schismatosphaeridium</i> sp. B *																			
<i>Pterospermella hermosita</i>																			
<i>Polyedryxium</i> cf. <i>P. decorum</i>																			
<i>Estiastra culcita</i>																			
<i>Polyedryxium simplex</i>																			
<i>Bimerga paulae</i>																			
<i>Cordobesia orientalis</i>																			
<i>Cymatiosphaera salopensis</i>																			
<i>Dupliciradiatum tenue</i>																			
<i>Estiastra</i> sp. -																			
<i>Hoegklingia</i> cf. <i>H. longispina</i>																			
cf. <i>Visbysphaera</i> sp. A +																			
<i>Baculatretilatus</i> sp.																			
<i>Multiplicisphaeridium</i> spp.																			
<i>Estiastra</i> spp.																			
cf. <i>Riculusphaera fissa</i>																			
<i>Polyedryxium</i> cf. <i>P. condensum</i>																			
<i>Polyedryxium embudum</i>																			
<i>Stellinium rabians</i>																			
<i>Polyedryxium evolutum</i>																			
<i>Cymatiosphaera</i> aff. <i>C. ledburica</i> °																			
<i>Cymatiosphaera mirabilis</i>																			
<i>Polyedryxium robustum</i>																			
<i>Polyedryxium helenaster</i>																			
? <i>Polyedryxium asperum</i>																			
<i>Ammonidium</i> sp.																			
<i>Cymatiosphaera multisepta</i>																			
<i>Cymatiosphaera prismatica</i>																			
<i>Dictyotidium dictyotum</i>																			
cf. <i>Duvernaysphaera wilsonii</i>																			
<i>Leiofusa berneseae</i>																			
<i>Multiplicisphaeridium</i> cf. <i>M. rochesterense</i>																			
<i>Navifusa bacilla</i>																			
<i>Neoverhachium carminae</i>																			
<i>Palacanthus ledanoisii</i>																			
<i>Villosacapsula setosapellucula</i>																			
<i>Palacanthus stelligerum</i>																			
<i>Veryhachium trispiniflatum</i>																			

Phytoplankton









Highlights

- Devonian palynological assemblages from the Argentinian Precordillera are presented
- Marine and terrestrial palynomorphs suggest a late Pragian to Emsian age
- The age of the guide horizon of Keidel is constrained to the late Pragian
- Palynological data enable a new biostratigraphic frame for the Devonian units