

1 **SILURIAN CONODONTS FROM THE RINCONADA FORMATION,**
2 **ARGENTINE PRECORDILLERA**

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24 **Abstract.** At the eastern margin of the Precordillera and close to the boundary with the
25 Sierras Pampeanas, the Rinconada Formation displays extensional faults, boudinaged
26 sequences and scarce slump folds whose age are not well constrained. It represents a stacking
27 of “broken formations” intercalated with scarce debris flows. A calcareous iron-rich sample
28 from the upper levels of the Rinconada Formation yielded 134 conodont elements including
29 *Dapsilodus obliquicostatus*, *Decoriconus fragilis*, *Oulodus* sp., *Pseudooneotodus*
30 *beckmanni*, *P. b. bicornis*, *Wurmiella excavata* along with ozarkodinids of the “Oz.”
31 *bohémica* – “Oz.” *snajdri* lineage. The record of “Oz.” aff. *snajdri* suggests a late Homeric
32 – early Gorstian (late Wenlock – early Ludlow) minimum depositional age for the upper
33 debris flows of the Rinconada Formation. The studied conodont fauna improves the
34 biostratigraphic correlation of the Rinconada melange to other successions in the
35 Precordillera and the world as well as it provides clues on the tectonostratigraphic
36 development of the SW Gondwanan foreland during the Silurian times.

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38 **Keywords:** conodonts, biostratigraphy, Rinconada Formation, Silurian, Precordillera,
39 Argentina.

40

41 **Resumen.** CONODONTES SILÚRICOS DE LA FORMACIÓN RINCONADA,
42 PRECORDILLERA ARGENTINA. La Formación Rinconada, situada en el margen oriental
43 de la Precordillera y próxima al límite con las Sierras Pampeanas, muestra fallas
44 extensionales, así como pliegues y budines originados por deformación de sedimentos no
45 litificados. Esta formación, cuya edad no ha sido bien precisada, está constituida por un
46 apilamiento de “formaciones rotas” intercaladas con escasos flujos de detritos. Una muestra
47 calcárea rica en hierro de los niveles superiores de la Formación Rinconada brindó 134

48 conodontes, incluyendo *Dapsilodus obliquicostatus*, *Decoriconus fragilis*, *Oulodus* sp.,
49 *Pseudooneotodus beckmanni*, *P. b. bicornis*, *Wurmiella excavata* y ozarkodínidos del linaje
50 “Oz.” *bohémica* – “Oz.” *snajdri*. El registro de “Oz.” aff. *snajdri* permite sugerir una edad
51 mínima depositacional Homeriana tardía – Gorstiana temprana (Wenlock tardío – Ludlow
52 temprano) para los flujos de detritos del tramo superior de la Formación Rinconada. La fauna
53 de conodontes recuperada de la Formación Rinconada favorece la correlación estratigráfica
54 con otras sucesiones sedimentarias de la Precordillera y del mundo, así como brinda claves
55 sobre el desarrollo tectonoestratigráfico del antepaís del suroeste de Gondwana en tiempos
56 silúricos.

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58 **Palabras clave:** conodontes, bioestratigrafía, Formación Rinconada, Precordillera,

59 Argentina.

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72 **INTRODUCTION**

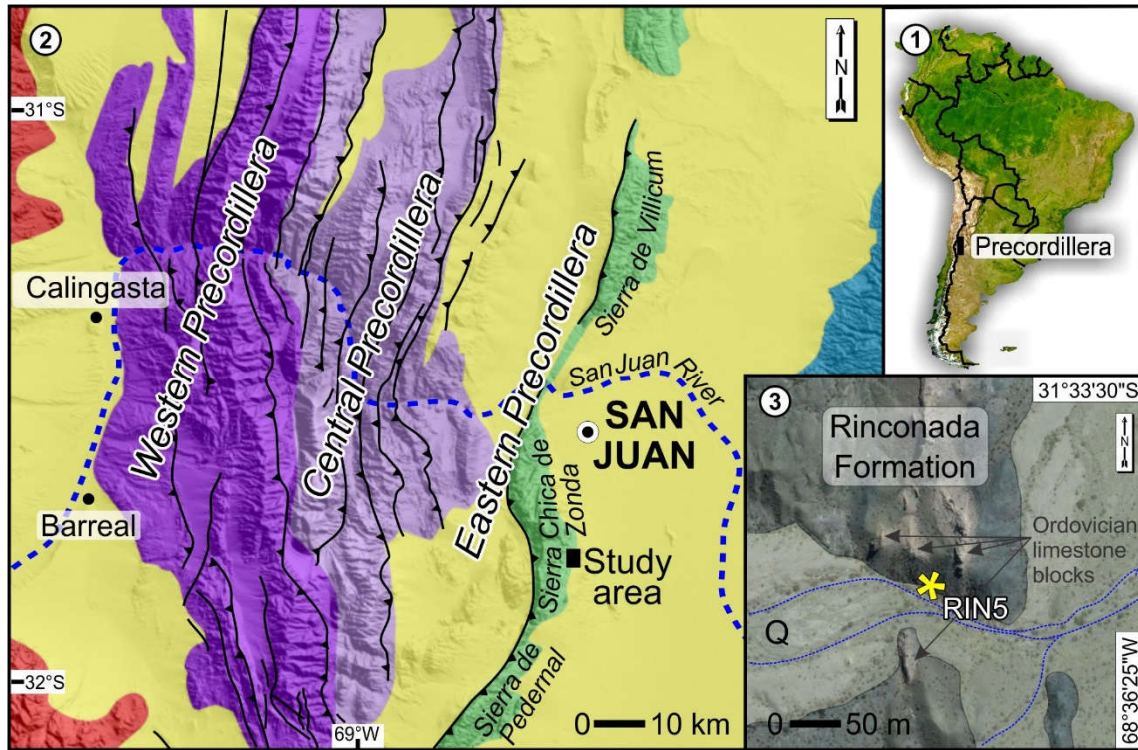
73 In the eastern margin of the Argentine Precordillera, the Rinconada melange (Amos, 1954;
74 Gosen *et al.*, 1995) accounts for a major rearrangement process in the early Palaeozoic
75 basin nearby the boundary with Sierras Pampeanas. The genesis and age of this melange
76 are controversial due to the inherent reworked character of its components but they are
77 essential to decipher the interrelationship between both continental blocks during the
78 building of southwestern Gondwana (*e.g.*, Gosen *et al.*, 1995; Ramos *et al.*, 1996; Rapela *et*
79 *al.*, 2016).

80 After a multidisciplinary study of the Rinconada Formation committed to decipher its
81 geological significance, we conducted a systematic conodont sampling along with detailed
82 structural and sedimentological analysis. We processed 46 rock samples (85 kg total
83 weight) from carbonate-cemented sandstones, conglomerates and blocks included in the
84 melange, which yielded Early Ordovician, Middle Ordovician (Voldman *et al.*, 2015), and
85 a unique Silurian conodont fauna that is subject of this contribution.

86 The Silurian conodont collection proceeds from a calcareous sandstone incorporated in a
87 debris flow from the upper levels of the Rinconada Formation (Figures 1, 2). The
88 preservation of the specimens is in general poor, probably related to the slumping forces
89 that gave rise to the melange, but provided significant biostratigraphic information. In
90 addition to the long-range simple cones *Decoriconus*, *Dapsilodus* and *Pseudooneotodus*,
91 the conodont collection includes specimens of the genera *Oulodus*, “*Ozarkodina*” and
92 *Wurmiella*, which suggests a late Silurian age for the upper levels of the Rinconada
93 Formation. The purpose of the present contribution is to enhance the biostratigraphic
94 correlation of the Rinconada Formation to other successions in the Precordillera and basins

95 of the world, as well as shed light on the scarcely known Silurian conodont faunas from
96 SW Gondwana.

97 [Figure 1]



98
99 **Figure 1.1.** Geographic location of the Precordillera, Argentina; **2,** Location map of the
100 Rinconada Formation study area, on the eastern flank of the Sierra Chica de Zonda, Eastern
101 Precordillera; **3,** Geographic location of the fossiliferous sample RIN5, Rinconada
102 Formation, San Juan Province.

103 **GEOLOGICAL SETTING**

104 The Rinconada Formation is a melange located in the Eastern Precordillera, a west-verging
105 imbricate thrust system developed in Silurian times (*e.g.*, Ramos *et al.*, 1996). The
106 Rinconada melange crops out along the eastern slopes of Sierra Chica de Zonda, Sierra de
107 Pedernal (Fig. 1), and also in the Sierra de Villicum, where it has been named Mogotes
108 Negros Formation (Cuerda, 1985; Benedetto *et al.*, 1992; Peralta, 1993). Its geological

109 significance is controversial, being considered either as the result of tectonic (Keidel, 1938;
110 Heim, 1948) or gravity-driven processes (Amos, 1954; Borrello, 1969; Harrington, 1971;
111 Amos and Fernandez, 1977; Peralta, 1990; Gosen *et al.*, 1995).

112 First fossil findings from the Rinconada Formation were reported by Keidel (1938), who
113 identified *Atrypina acutiplicata* (= *Harringtonina australis*) and *Monograptus* sp. at its type
114 locality (La Rinconada). Amos and Fernández (1977) reported the brachiopod *Leptocoelia*
115 *nunezi* (Early Devonian) from an equivalent sequence at Cerro Bola, to the south of the
116 type section. Cuerda (1981) described from the upper levels of the Rinconada Formation a
117 graptolitic association composed of *Climacograptus* cf. *minutus*, *Diplograptus* sp. and
118 *Monograptus* sp., which referred to the Llandovery, supporting the early ideas of Keidel
119 (1938) and Heim (1948), who postulated a Silurian age for all or at least parts of the
120 Rinconada Formation. Benedetto and Franciosi (1998) suggested that the uppermost strata
121 of the Rinconada Formation may be older than late Wenlock based on the record of the
122 inarticulate brachiopod *Leangella* (*Leangella*) in the Sierra Chica de Zonda.

123 The age of the base of the Rinconada Formation is less constrained. For instance, Peralta
124 and Uliarte (1986) recovered middle Darriwilian graptolites from the lower levels of the
125 Rinconada Formation, which they interpreted as deposited over the San Juan Formation in
126 transitional contact. Sarmiento *et al.* (1988) studied a conodont collection from correlative
127 levels and supported a middle Darriwilian age for the top of the San Juan Formation and the
128 base of the Rinconada Formation though the samples from the latter unit correspond to
129 allochthonous blocks. Indeed, carbonate blocks of decametric size as well as limestone
130 clasts incorporated in polymictic conglomerates yielded Early to Middle Ordovician
131 conodont specimens as recognized in recent studies (Lehnert, 1995; Voldman *et al.*, 2015).

132 In the Don Braulio section, situated on the eastern flank of Sierra de Villicum, the

133 Rinconada Formation overlies the Don Braulio Formation, which bears graptolites of
134 Llandovery age (Peralta, 1986). Gosen *et al.* (1995) suggested that the Rinconada
135 Formation is Silurian but did not discard an Ordovician age for parts within or at the base
136 of the melange.

137 In the Rinconada locality (Sierra de Chica de Zonda), the melange is at least 2500 m-thick
138 and overlies limestones of the San Juan Formation. It is composed mainly of sandstone-
139 shale alternations, massive shales, conglomerates and limestone boulders up to km-scale in
140 length. In most places, these lithostratigraphic units display ubiquitous extensional faults
141 with variable extension values. There are also boudinaged sequences and scarce slump
142 folds, which record deformation of not well-lithified sediments. Therefore, most of the
143 Rinconada melange could be regarded as a stacking of “broken formations” *sensu*
144 Raymond (1984), intercalated with scarce debris flows that mainly occur in the upper part
145 of the succession. The arrangement of the deposits suggests submarine mass-transport
146 processes probably at the orogenic wedge front (for a review see Festa *et al.*, 2016).

147 From the upper levels of the sedimentary succession comes the conodont sample RIN 5
148 (31°44'39.78"S, 68°36'35.72"W, Figures 1, 2), which is a lenticular reddish calcareous
149 sandstone with iron oolites, similar to the fossiliferous samples studied by Benedetto and
150 Franciosi (1998). Successively, the Rinconada Formation is covered through angular
151 unconformities by Carboniferous and Neogene sedimentary rocks (Heim, 1948; Amos,
152 1954), which are involved in thrusting and folding related to the Andean orogeny.

153 **[Figure 2]**



154

155 **Figure 2.1.** Limestone blocks from the San Juan Formation embedded in debris flows from
156 the upper levels of the Rinconada Formation, proximate to the Neogene unconformity
157 (view to the north); **2.** Iron-rich calcareous level Rin5, which provided the first Silurian
158 conodont fauna from the Rinconada Formation (for location see figure 1); **3.** Debris flow
159 from the upper levels of the Rinconada Formation showing a cleaved matrix with pelitic,
160 siliceous and calcareous intraclasts. Hammer for scale is 33 cm long.

161 **METHODS**

162 The Silurian conodont collection consists of 134 elements (and tens of fragments) that were
163 obtained after digestion of 4.25 kg of rock in 10% formic acid solution, following the
164 standard acid etching techniques described by Stone (1987). The conodont elements appear
165 frequently broken and show a conodont Colour Alteration Index (CAI) 3, which indicates
166 burial palaeotemperatures ranging from 110–200 °C (Epstein *et al.*, 1977). The abundant
167 microfractures of the specimens inhibited ultrasonic cleaning, as test specimens resulted
168 severely damaged. Conodont images were acquired with an Olympus LEXT OLS4000
169 confocal microscope at the LAMARX (Universidad de Córdoba) and with a Leica M205FA
170 Microscope of the Servicios Científico-Técnicos (Universidad de Oviedo). The conodont
171 elements are housed under repository code CORD-MP 50815 up to 50949 in the Museo de
172 Paleontología, Facultad de Ciencias Exactas, Físicas y Naturales, Universidad Nacional de
173 Córdoba, Argentina.

174 **SYSTEMATIC PALAEOLOGY**

175 Class CONODONTA Pander, 1856

176 Order BELODELLIDA Sweet, 1988

177 Family DAPSILODONTIDAE Sweet, 1988

178 Genus *Dapsilodus* Cooper, 1976

179 **Type species.** *Distacodus obliquicostatus* Branson and Mehl, 1933.

180 **Discussion.** *Dapsilodus* has an apparatus composed of laterally compressed coniform
181 elements characterized by distinctive chevron-shaped ornamentation adjacent to the
182 anterior margin (Cooper, 1976; Barrick, 1977). It includes M elements (acodontiform) and
183 a symmetry transition series. The Sa elements (modified distacodontiform) are
184 symmetrical, have a long base, and a lateral costa on each side that may extend behind the
185 posterior keel. The Sb elements (distacodiform) are slightly twisted whereas the Sc
186 elements (distacodiform) are strongly twisted. The Sb-Sc elements in *Dapsilodus* are
187 morphologically similar to their homologous of the Ordovician genus *Besselodus*.
188 However, in the latter genus the distacodiform elements are straight, not twisted, and the M
189 elements are oistodiform (*e.g.*, Nowlan *et al.*, 1988).

190 ***Dapsilodus obliquicostatus*** (Branson and Mehl, 1933)

191 Figure 3.12

192 1933. *Distacodus obliquicostatus*. Branson and Mehl, p. 41, pl. 3, fig. 2.

193 1978. *Distacodus obliquicostatus* Branson and Mehl. Rexroad *et al.*, p. 4, pl. 1, fig. 9.

194 1991. *Distacodus obliquicostatus* Branson and Mehl. McCracken, p. 78–79, pl. 4, figs. 11,
195 13–14, 16–28, 30–32, 35, 40. (*cum. syn.*).

196 1999. *Dapsilodus obliquicostatus* (Branson and Mehl). Lehnert *et al.*, pl. 1, figs. 1–8.

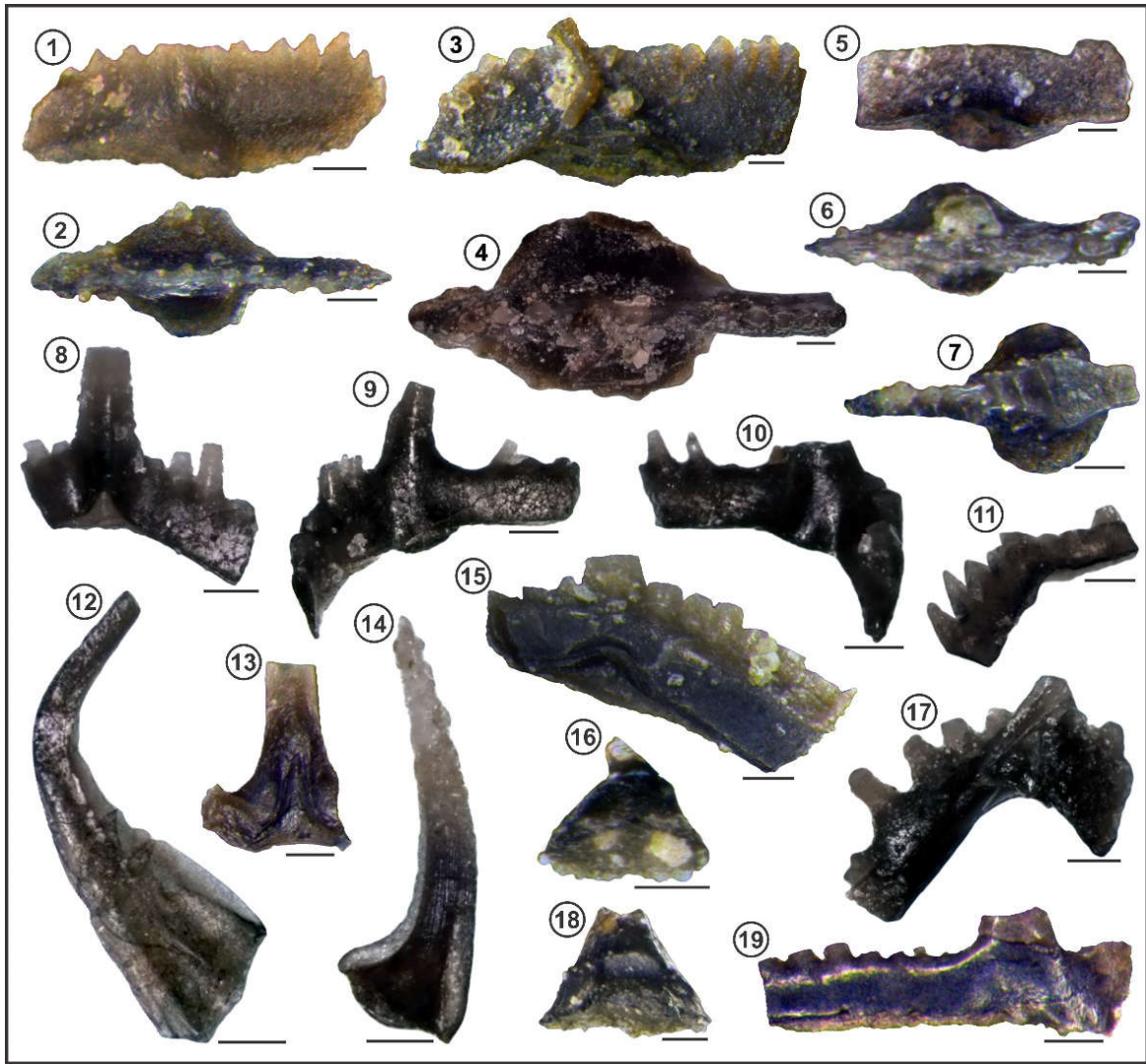
197 1999. *Dapsilodus obliquicostatus* (Branson and Mehl). Cockle, p. 119, pl. 4, figs. 13–14,
198 17–19 (non 15–16).

199 2009. *Dapsilodus obliquicostatus* (Branson and Mehl). Mestre, p. 474, 476, figs. 3.12–3.15.
200 **Material.** 3 Sa, 16 Sb, 16 Sc elements (CORD-MP 50815-50849).

201 **Discussion.** The studied specimens display the typical oblique striations of the genus in the
202 antero-basal margin. The base is broad and long, frequently extended to posterior. The
203 basal cavity is deep and tapers gradually to the point of maximum curvature, with the apex
204 centrally located. The anterior and posterior margins are keeled along the entire length of
205 the elements. Soft lateral costae are present near midface. M elements were not recovered.

206 *Dapsilodus obliquicostatus* is a widespread and long-ranging Silurian taxon that inhabited
207 mainly off-shore to oceanic environments (*e.g.*, Aldridge and Jeppsson, 1984; Armstrong,
208 1990; McCracken, 1991; Zhang *et al.*, 2006). Aldridge and Mabillard (1981) reported a
209 strikingly high proportion of *D. obliquicostatus* in the furthest offshore localities across the
210 Early Silurian shelf of the Wales Borderland area. It is also present in Silurian deposits of
211 Australia (Cockle, 1999), Austria (Suttner, 2007), China (Lin, 1983), Czech Republic
212 (Slavík, 2014), Greenland (Armstrong, 1990), Sweden (Bergström and Bergström, 1996;
213 Dahlqvist and Bergström, 2005), Morocco (Benfrika *et al.*, 2007; Corrigan *et al.*, 2014),
214 North America (Cooper, 1976; McCracken, 1991), Russia (Zhang and Barnes, 2007; Kaljo
215 *et al.*, 2012), and Spain (Martín Algarra *et al.*, 2009), among other places.

216 **[Figure 3]**



217

218 **Figure 3.** Silurian conodonts from the Rinconada Formation, sample RIN5, Precordillera
 219 Argentina. 1–4, *“Ozarkodina”* aff. *snajdri* (Walliser), 1–2, Pa element, lateral and oral
 220 view, CORD-MP 50913; 3–4, Pa element, lateral and oral views, see attached coniform
 221 element on lateral side, CORD-MP 50914. 5–6, *“Ozarkodina”* sp., Pa element, CORD-MP
 222 50918, lateral and oral views. 7–8, 11, undetermined "ozarkodinids", 7, Pa element,
 223 CORD-MP 50941, oral view; 8, Sa element, CORD-MP 50946, dorsal view; 11, Pb
 224 element, CORD-MP50948, lateral view. 9–10, 17, *Oulodus* sp., 9, Sb element, CORD-MP
 225 50920, lateral view; 10, Sb element, CORD-MP 50922, lateral view; 17, P element, CORD-

226 MP 50943, dorsal view. **12**, *Dapsilodus obliquicostatus* (Branson and Mehl), Sc element,
227 CORD-MP 50815, lateral view. **13**, **15**, **19**, *Wurmiella excavata* (Branson and Mehl), **13**,
228 Sa element, CORD-MP50921, dorsal (posterior) view; **15**, Pa element, CORD-MP 50919,
229 lateral view; **19**, Sc element, CORD-MP 50923, lateral view. **14**, *Decoriconus fragilis*
230 (Branson and Mehl, 1933), Sb element, CORD-MP 50850, lateral view. **16**,
231 *Pseudooneotodus beckmanni* (Bischoff and Sannemann), P element, CORD-MP 50873,
232 lateral view. **18**, *Pseudooneotodus bicornis bicornis* (Drygant), P element, CORD-MP
233 50870, lateral view. Scale bar: 0.1 mm.

234 Order Prioniodontida Donoghue *et al.*, 2008

235 Suborder Ozarkodinida Donoghue *et al.*, 2008

236 **Remarks.** Murphy *et al.* (2004) provided a major taxonomical revision of late Silurian and
237 Early Devonian ozarkodinids. They introduced *Wurmiella* to house taxa related to the
238 “*excavata* Group”, *Zieglerodina* to include the ozarkodinids of the “*remscheidensis* Group”,
239 and an informal genus (“Genus W”) to contain the “*eosteinhornensis* stock”. Their proposal
240 was rejected by Suttner (2007, 2009), but adopted with some modifications by others (*e.g.*,
241 Carls *et al.*, 2007; Corrigan and Corradini, 2009; Drygant and Szaniawski, 2012; Corrigan *et*
242 *al.*, 2014; Jarochowska and Munnecke, 2015). Donoghue *et al.* (2008) verified the
243 polyphyletic character of the multiple species assigned to *Ozarkodina*, and pointed out that
244 “*Nicollidina* (= *Zieglerodina*) *remscheidensis* (Ziegler, 1960)” is closely related to the
245 Frasnian polygnathid *Nicollidina brevis* Dzik (2002). On the other hand, “*Oz.*” *bohémica*
246 (Walliser, 1964), “*Oz.*” *crispa*, “*Oz.*” *sagitta rhenana* (Walliser, 1964), and “*Oz.*” *s. sagitta*
247 (Walliser, 1964), would conform a ‘stem’ group along with *Oz.*” *snajdri* and “*Oz.*”
248 *parasnajdri* (Walliser, 1964), given the marked morphological differences of their Pa

249 elements (*e.g.*, lack of anterior cockscomb, far posterior extension of basal lobes, fused
250 denticles) when compared to *Ozarkodina* s.s. (*e.g.*, Slavík and Carls, 2012; Peavey, 2015).

251 **“*Ozarkodina*” aff. *snajdri*** (Walliser, 1964)

252 Figures 3.1–3.4, 3.6, 4.1–4.4

253 **Material.** 5 Pa elements (CORD-MP 50913-50917).

254 **Remarks.** The Pa elements attributed to “*Ozarkodina*” aff. *snajdri* include transitional forms
255 of the “*Oz.*” *bohemica*–*snajdri* group (Figs. 3.1–3.4, 4.1–4.4). They have denticles of
256 subequal size, stout, fused except for discrete tips, and slightly tilted ventrally. Denticles
257 located above the aboral cavity are completely fused so conform a crenulated ridge. The
258 platform is spear-shaped, unornamented and weakly asymmetrical, with the halves on both
259 sides of the blade slightly offset from each other. It extends approximately from the centre
260 of the blade to just before its dorsal edge. The basal margin profile is stepped, with the
261 anterior section slightly bowed. The anterior process has a steeply curved termination, with
262 a distal denticle approximately half way up blade (Figs. 3.1, 3.3). The posterior process also
263 reduces rapidly in size at its end. White matter concentrates at the tip of the denticles and
264 gradually diffuses toward the blade whereas the basal cavity tip is not discernible due to the
265 opaque aspect of the elements.

266 Helfrich (1975) introduced *Spathognathus tillmani* from the lower Wills Creek Formation,
267 central Appalachian Mountains, to include specimens grossly similar to *S. sagitta bohemicus*
268 and *S. snajdri*, but with a slightly longer blade and the basal platform more posteriorly
269 located. Helfrich's illustrations of “*Oz. tillmani*” include Pa elements very similar to elements
270 of this study, but differ in the more developed denticulation. “*Oz. tillmani*” was subsequently
271 considered a *nomem dubium* because the systematics appeared on microfiche (Cooper, 1980)
272 and regarded as a junior synonym of “*Oz.*” *bohemica* (Calner and Jeppson, 2003). Currently,

273 “*Oz.*” *bohemica* s.l. encompass a group of forms with considerable variability, even within
274 collections, that are frequently described as morphotypes or subspecies (Schönlaub in Kříž
275 *et al.*, 1993; Viira and Aldridge, 1998; Calner and Jeppson, 2003).

276 In particular, “*Oz.*” aff. *snajdri* is distinguished from typical representatives of “*Oz.*” *snajdri*
277 by having a blade of more uniform height and a less expanded, more symmetrical, and
278 centrally located basal cavity. Notwithstanding that, specimens attributed to “*Oz.*” *snajdri*
279 with a relatively small basal cavity and a straight upper margin are included in the literature
280 (e.g., Miller, 1995; Viira and Aldridge, 1998). “*Oz.*” aff. *snajdri* also shows resemblance to
281 *Oz. bohemica* morphotype 3 of Schönlaub and the coeval *Oz. bohemica longa* (Calner and
282 Jeppson, 2003; Slavík, 2014), but differ in the more centrally located basal expansion, the
283 gently convex anterior and posterior edges, and the shorter blade length. On the other hand,
284 *Oz. inflata* (Walliser, 1964) is distinct as it possesses the oral margin convex, lacks denticle
285 fusion, and the basal lobes are more symmetrical. The transitional character of the “*Oz.*”
286 *snajdri* lineage and the scarce number of specimens recovered from the Rinconada Formation
287 inhibits a further refinement of the taxonomic designation. Eventually, more records of “*Oz.*”
288 *tillmani* and of “*Oz.*” aff. *snajdri* would help determine if they are conspecific.

289 **“*Ozarkodina*” sp.**

290 Figures 3.5–3.6, 4.5–4.6

291 **Material.** 1 Pa element (CORD-MP 50918).

292 **Remarks.** “*Ozarkodina*” sp. is characterized by having a small basal cavity relatively
293 centrally located and the oral margin adenticulated. The basal cavity lips prolong up to the
294 posterior tip of the blade, whereas they continue as a narrow groove towards the anterior
295 edge. The anterior and posterior margins of the blade are fairly straight.

296 “*Ozarkodina*” sp. resembles “*Oz.*” *parasnajdri* (Viira and Aldridge, 1998) in the
297 adenticulated oral margin and in the basal cavity position, though it lacks the large basal
298 cavity that characterizes the representatives of the “*Oz.*” *snajdri*–*parasnajdri* group. It is
299 also morphologically similar to “*Oz.*” *b. longa*, but it has the basal cavity more centrally
300 located. Moreover, in “*Oz.*” *b. longa* the fused ridges of denticles occur mainly in mature to
301 gerontic elements, though they are more frequent in the younger populations (Calner and
302 Jeppsson, 2003).

303 Genus *Wurmiella* Murphy *et al.*, 2004

304 **Type species.** *Ozarkodina excavata* subspecies *tuma* Murphy and Matti, 1983

305 ***Wurmiella excavata*** (Branson and Mehl, 1933)

306 Figures 3.13, 3.15, 3.19, 4.8, 4.9

307 1933. *Prioniodus excavatus* Branson and Mehl, p. 45, pl. 3, figs. 7, 8.

308 2005. *Ozarkodina excavata* (Branson and Mehl). Bitter and Purnell, text-figs. 2–3, pl. 1–3.

309 2006. *Ozarkodina excavata excavata* (Branson and Mehl). Albanesi *et al.*, pl. 5, figs. M–Y.

310 2009. *Wurmiella* cf. *excavata* Branson and Mehl. Mestre, p. 474, fig. 3: 4–6.

311 2016. *Wurmiella excavata* (Branson and Mehl 1933). Sullivan *et al.*, fig. 4: 3–12.

312 2016. *Ozarkodina excavata* (Branson and Mehl). Mathieson *et al.*, p. 624, 626, figs 20,

313 30W, X, 32L–Q (*cum. syn.*).

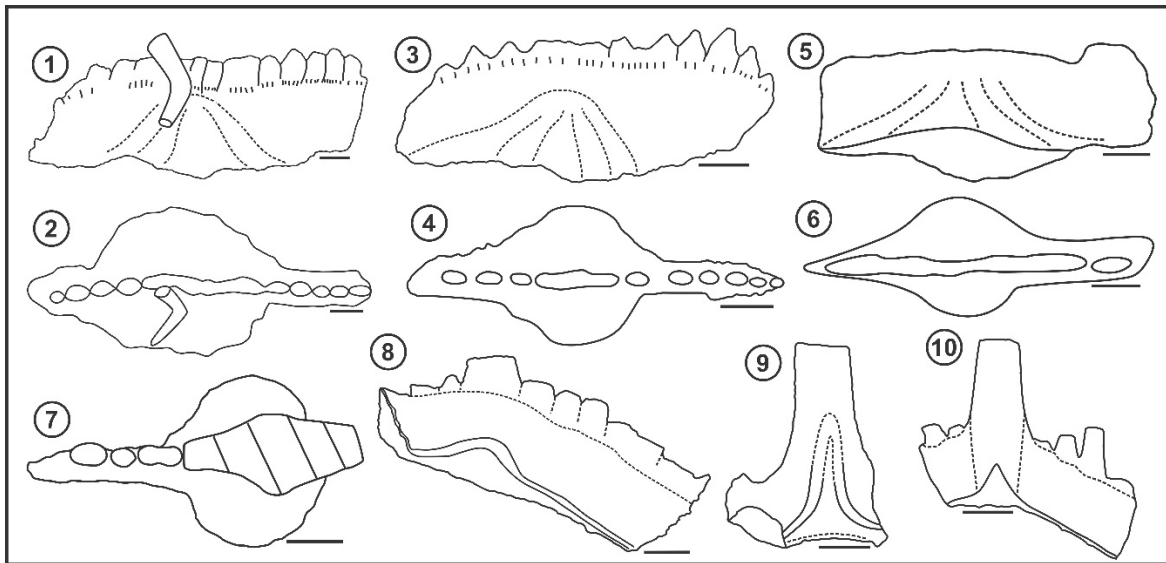
314 **Material.** 17 Pa, 1 Pb, 1 Sa, 2 Sb, 1 Sc elements (CORD-MP 50919-50940).

315 **Discussion.** *Wurmiella excavata* is frequently reported in many Silurian – Lower Devonian
316 collections, yet there is a large uncertainty surrounding its type specimens (Murphy *et al.*,
317 2004). In addition, it exhibits significant morphological variations through time probably
318 associated with variations in diet (Jones and Purnell, 2007). Donoghue *et al.* (2008) suggested
319 that *Wurmiella* may not be its appropriate taxonomic home, but awaits further phylogenetic

320 evidence to establish it. The specimens from the Rinconada Formation fit well with respect
321 to the typical material from the Bainbridge Limestone (Branson and Mehl 1933).

322 *Wurmiella excavata* is a very successful component of conodont communities, showing large
323 resistance against palaeoenvironmental changes. It has a wide paleobiogeographic
324 distribution, with records in the Carnic Alps (Walliser, 1964; Corradini *et al.*, 2015), North
325 America (Branson and Mehl, 1933; von Bitter and Purnell, 2005), Northern Australia
326 (Simpson, 2000), Bohemia (Slavík *et al.*, 2010), Central Iran (Männik *et al.*, 2013), as well
327 as in the Los Espejos Formation (Albanesi *et al.*, 2006) and the Tambolar Formation (Mestre,
328 2009) in the Precordillera.

329 **[Figure 4]**



330

331 **Figure 4.** Line-art drawings of selected specimens from sample RIN5. **1–4, “Ozarkodina”**
332 **aff. *snajdri*, 1–2, Pa element, lateral and oral view, CORD-MP 50914; 3–4, Pa element,**
333 **lateral and oral views, see attached coniform element on lateral side, CORD-MP 50913. 5–**
334 **6, “Ozarkodina” sp., Pa element, CORD-MP 50918, lateral and oral views. 7, undetermined**
335 **“ozarkodinid”, Pa element, CORD-MP 50941, oral view. 8–9, *Wurmiella excavata* (Branson**

336 and Mehl), **8**, Pa element, CORD-MP 50919, lateral view; **9**, Sa element, CORD-MP50921,
337 dorsal (posterior) view. **10**, undetermined "ozarkodinid", Sa element, CORD-MP 50946,
338 dorsal view. Scale bar: 0.1 mm.

339 Order Prioniodinina Donoghue *et al.*, 2008

340 Family PRIONIODINIDAE Bassler, 1925

341 Genus *Oulodus* Branson and Mehl, 1933

342 **Type species.** *Cordylodus serratus* Stauffer, 1930

343 *Oulodus* sp.

344 Figures 3.9, 3.10, 3.17

345 **Material.** 1 P, 2 Sb, 1 S (CORD-MP 50943-50945).

346 **Discussion.** We follow here the diagnosis of *Oulodus* given by Sweet and Schönlaub (1975),
347 which implies a conservative apparatus composition through the Middle Ordovician and
348 Devonian. Slavík *et al.* (2010) adopted the genus name *Delotaxis* rather than *Oulodus* for the
349 late Silurian-early Devonian taxa with digyrate apparatuses and five pairs of different
350 elements. However, the upper Silurian species have seximembrate apparatuses, so belong to
351 *Oulodus* (C. Corradini, written com.). Besides, Donoghue *et al.* (2008) observed that a
352 definition of sexi- and septimembrate apparatus in Prioniodinina is meaningless due to the
353 architectural stability of the apparatus within Ozarkodinida. Thus, the number of elements in
354 the apparatus may reflect the subjective assessment of the morphological boundaries between
355 element morphotypes.

356 The collection from the Rinconada Formation includes rather few fragmented elements that
357 complicate the specific determination. These exhibit subcircular cusps and discrete denticles
358 of similar height and round cross sections consistent with the morphological characteristics

359 of *Oulodus*. The record of two Sb elements with the anterior process with different angles of
360 curvature imply, eventually, the presence of two different species.

361 Order ?PROTOPANDERODONTIDA Sweet, 1988

362 Family ?PROTOPANDERODONTIDAE Lindström, 1970

363 Genus *Decoriconus* Cooper, 1975

364 **Type species.** *Paltodus costulatus* Rexroad, 1967.

365 **Discussion.** *Decoriconus* includes small, generally striated and laterally compressed
366 coniform elements. The lineage can be traced back up to the Tremadocian with *D.*
367 *peselephantis* (Lindström) and it extends up to the Devonian with *D. fragilis* (e.g., Löfgren,
368 1998; Drygantand Szaniawski, 2012). Cusps and keels are albid whereas hyaline material
369 surrounds the basal cavity. The abrupt transition between hyaline and albid material conforms
370 a straight line that meets the anterior margin of the cusp at an angle of *ca.* 90° in Ordovician
371 elements, which reduces to *ca.* 60° in the Silurian specimens (Löfgren, 1998).

372 ***Decoriconus fragilis*** (Branson and Mehl, 1933)

373 Figure 3.14

374 1933. *Paltodus fragilis* Branson and Mehl, p. 43, pl. 3, fig. 6.

375 1978. *Decoriconus? fragilis* (Branson and Mehl). Rexroad *et al.*, p. 4, pl. 1, fig. 10.

376 1991. *Decoriconus fragilis* (Branson and Mehl). McCracken, p. 79–80, pl. 4, figs. 33, 39.

377 2002. *Decoriconus fragilis* (Branson and Mehl). Zhang and Barnes, p. 11–12, gs. 15.25–
378 15.31.

379 2007. *Decoriconus fragilis* (Branson and Mehl). Zhang and Barnes, p. 505, figs. 9.10–9.15.

380 2007. *Decoriconus fragilis* (Branson and Mehl). Suttner, p. 19–20, pl. 7, figs. 12–21 (*cum.*
381 *syn.*).

382 2015. *Decoriconus fragilis* (Branson and Mehl). Jarochowska and Munnecke, fig. 7: O–S,
383 fig. 8: M–Q.

384 **Material.** 1 Sa, 18 Sb, 1 Sc elements (CORD-MP50850-50869).

385 **Discussion.** Branson and Mehl (1933) defined *Paltodus fragilis* with specimens recovered
386 from shales of the Bainbridge Formation in southeastern Missouri. Cooper (1976)
387 reconstructed the oral apparatus of *D. fragilis* including acontiodontiform, paltodontiform
388 and drepanodontiform elements. Barrick (1977) referred to them as Sa, Sb, and Sc elements,
389 respectively, because they bear little resemblance to the shapes represented by the original
390 genera, following the alphabetic locational scheme of Sweet and Schönlaub (1975).
391 McCracken and Barnes (1981) included an Sc element in the apparatus reconstruction of *D.*
392 *costulatus*, whose lack was previously attributed to a different apparatus composition. Based
393 on this record, Zhang and Barnes (2002) suggested that *D. costulatus* is a junior synonym of
394 *D. fragilis*.

395 According to the description of Barrick (1977), the Sa element is strongly compressed, nearly
396 symmetrical and slightly twisted. The anterior margin is keeled and merges smoothly into
397 the lateral faces. The posterior margin is also keeled but surrounded by a lateral groove on
398 each side. The Sb element is asymmetrical, twisted inwardly and recurved near midheight. It
399 has a moderately flaring base and sharp anterior and posterior keels. The latter is separated
400 from the lateral faces by narrow grooves. The distinct Sc element is sharply incurved, with
401 the anterior margin of the base extended as an anticusp of variable lengths.

402 The collection *D. fragilis* from the Rinconada Formation include Sa and Sb elements with
403 well-developed keels on both anterior and posterior margins, even on the base, associated
404 with relatively shallow grooves. In contrast, the recovered Sc element presents the anterior

405 margin keeled and the posterior one subrounded, which is consistent with some previous
406 reports (e.g., Kleffner, 1987, fig. 6.20).

407 *Decoriconus fragilis* is common in outer shelf to basin environments, in some cases more
408 than 100 m deep (e.g., Aldridge and Mabillard, 1981; Aldridge and Jeppsson, 1984;
409 Armstrong, 1990; Zhang and Barnes, 2002; Dahlgvist and Bergström, 2005). It is frequently
410 the dominant species along with *D. obliquicostatus* and *Aspelundia* in the Silurian outer shelf
411 and upper slope communities of Australia (Farrell, 2006), Austria (Suttner, 2007), Greenland
412 (Armstrong, 1990), North America (McCracken, 1991; Zhang and Barnes, 2002), Russia
413 (Zhang and Barnes, 2007) and Sweden (Bergström and Bergström, 1996).

414 Genus *Pseudooneotodus* Drygant, 1974

415 **Type species.** *Oneotodus? beckmanni* Bischoff and Sannemann, 1958

416 **Discussion.** *Pseudooneotodus* consists of short phosphatic cones arranged in an unimembrate
417 apparatus, whose suprageneric classification is not clearly defined (see Corradini, 2008, for
418 a recent discussion of the genus). It is made up of a lamellar cap underlain by a spherulitic
419 basal tissue characteristic of conodonts (Sanson, 1996). The different species of
420 *Pseudooneotodus* are classified according to the shape and number of apical tips and the
421 outline of the basal margin.

422 *Pseudooneotodus* is widely distributed in Middle Ordovician to Lower Devonian rocks all
423 over the world. In particular, Corradini (2008) observed that the occurrence of *Ps. beckmanni*
424 and *Ps. bicornis* is very irregular, even in successive beds in the same section, suggesting
425 that they were ecologically controlled to a higher degree than coeval conodont taxa. Both
426 species are typical from open-marine, offshore shelf environments.

427 ***Pseudooneotodus beckmanni*** (Bischoff and Sannemann, 1958)

428 Figure 3.16

429 1958. *Oneotodus? beckmanni* Bischoff and Sannemann, p. 98, pl. 15, figs. 22–25.
430 1988. *Pseudooneotodus beckmanni* (Bischoff and Sannemann). Hünicken and Sarmiento, p.
431 228, pl. 1, figs. 7a-b, pl. 3, figs. 10a-b.
432 2008. *Pseudooneotodus beckmanni* (Bischoff and Sannemann). Corradini, p. 142, pl. 1,
433 figs. 1-7.
434 2011. *Pseudooneotodus beckmanni* (Bischoff and Sannemann). Sumrall *et al.*, fig. 4: F.
435 2016. *Pseudooneotodus beckmanni* (Bischoff and Sannemann). Mathieson *et al.*, p. 597–
436 598, fig. 8: Q–S (*cum. syn.*).

437 **Material.** 40 P elements (CORD-MP 50873-50912).

438 **Discussion.** The squat, single apical, conical elements from the Rinconada Formation fit
439 well with the described and illustrated specimens of *Ps. beckmanni* by Corradini (2008),
440 who thoroughly analysed the species.

441 *Pseudooneotodus bicornis bicornis* (Drygant, 1974)

442 Figure 3.18

443 1974. *Pseudooneotodus bicornis* Drygant, p. 67, pl. 2, figs 40–48.

444 2008. *Pseudooneotodus bicornis bicornis* (Drygant). Corradini, p. 144, 146, pl. 1, figs. 8–
445 17.

446 2015. *Pseudooneotodus bicornis* Drygant, Jarochovska and Munnecke, fig. 7: J?

447 2016. *Pseudooneotodus bicornis* Drygant. Sullivan *et al.*, fig. 4: 15–16.

448 **Material.** 3 P elements (CORD-MP 50870-50872).

449 **Discussion.** Corradini (2008) split *Pseudooneotodus bicornis* in two subspecies, *Ps.*
450 *bicornis bicornis* and *Ps. bicornis contiguus*, based on the distance between the discrete
451 tips. The studied material consists of squat cones with subtriangular basal outline. It

452 exhibits two discrete tips in the apical part, separated by a depression, consistent with the
453 diagnosis of *Ps. bicornis bicornis*. (Corradini, 2008).

454 **BIOSTRATIGRAPHY**

455 In the absence of index fossils, the coniform specimens recovered from the Rinconada
456 Formation provide some general biostratigraphic information (Figure 5). For instance, *Ps.*
457 *beckmanni* ranges from the Ordovician up to the Emsian, but starts to occur in large numbers
458 in the Ludfordian, with two acme intervals during the Lochkovian and the Pragian, as
459 observed in Sardinia and the Carnic Alps (Corradini, 2008). *Ps. bicornis bicornis* is more
460 frequent during the upper Llandovery and Wenlock instead, uncommon in the Gorstian, and
461 it re-appears in the Ludfordian, with its LAD in the Lower *Oulodus elegans detortus* Zone
462 (Corradini, 2008). The species *D. obliquicostatus* extends for nearly the entire Silurian, while
463 its LAD marks the lower boundary of the Upper *Oulodus e. detortus* Zone in Sardinia and
464 the Carnic Alps (Corradini and Corrigan, 2012). Conversely, *D. fragilis* transits across the
465 terminal Ordovician mass extinction (Bergström *et al.*, 2012), ranging up to middle Pragian
466 in the Austrian Carnic Alps (Suttner, 2007).

467 **[Figure 5]**

AGE (Ma)				CONODONT ZONES					
	System	Series	Stage	Estonia	Carnic Alps	Prague Synform	Precordillera		
419.2	SILURIAN	PRIDOLI		<i>Oulodus elegans detortus</i>	Upper <i>Oulodus elegans detortus</i>				
							Lower <i>Oulodus elegans detortus</i>		
						<i>Ozarkodina r. remscheidensis</i>	<i>Ozarkodina eosteinhornensis</i> s.l. I.Z.		
						<i>Ozarkodina r. canadensis</i>			<i>Zieglerodina zellmeri</i>
						<i>Oz. r. eosteinhornensis</i>			
423.0				<i>Ozarkodina r. baccata / Oz. snajdri parasnajdri</i>			<i>Zieglerodina cf. zellmeri</i>		
		LUDLOW	Ludfordian	<i>Ozarkodina crispera</i>	<i>Ozarkodina crispera</i>	" <i>Ozarkodina</i> " <i>crispera</i>			
					<i>Ozarkodina snajdri</i>	<i>Pedavis latialata / Ozarkodina snajdri</i> I.Z.	" <i>Ozarkodina</i> " <i>parasnajdri</i> <i>Pedavis latialatus</i> <i>Parazieglerodina plodowskii</i> <i>Delotaxis fauna</i>		
					<i>Polygnathoides siluricus</i>	<i>Polygnathoides siluricus</i>	<i>Polygnathoides siluricus</i>		
					<i>Ancoradella ploeckensis</i>	<i>Ancoradella ploeckensis</i>	<i>Ancoradella ploeckensis</i>		
425.6					<i>Kockelella variabilis variabilis</i>	<i>Kockelella variabilis variabilis</i> I.Z.	<i>Kockelella variabilis variabilis</i> I.Z.	<i>Kockelella variabilis variabilis</i>	
427.4	WENLOCK	Homerian	<i>Kockelella crassa</i>	<i>Kockelella crassa</i>	<i>Kockelella crassa</i>				
				<i>Ctenognathus murchisoni</i>		<i>Kockelella ortus absidata</i>	"Oz." aff. <i>snajdri</i> ?		
				<i>Kockelella ortus absidata</i>		<i>Ozarkodina bohemica longa</i>			
				<i>Ozarkodina bohemica longa</i>		<i>Ozarkodina bohemica longa</i>			
				<i>Ozarkodina s. sagitta</i>	<i>Ozarkodina s. sagitta</i>	<i>Ozarkodina s. sagitta</i>			
430.5			<i>Kockelella ortus ortus</i>						

468
469 **Figure 5.** Comparison of late Wenlock – Pridoli conodont zones from Estonia (Männik,
470 2014), the Carnic Alps (Corradini *et al.*, 2015), the Prague Synform (Slavík and Carls, 2012;
471 Slavík, 2014), and the Precordillera (this study in grey background), with previous conodont
472 zones of Albanesi *et al.* (2006), Mestre (2009) and Sumrall *et al.* (2013).
473 The ramiform species *W. excavata* spans at least from the late Llandovery (*e.g.*, Cockle,
474 1999) up to the Emsian, with a wide paleobiogeographic distribution. It is frequently one of
475 the dominant species in the “Oz.” *b. longa*, *P. siluricus*, “Oz.” *crispera* and “Oz.”
476 *eosteinhornensis* zones (*e.g.*, Jeppsson, 1974; Mannik, 2007; Corrigan and Corradini 2009;
477 Fig. 5).
478 “Oz.” aff. *snajdri* offers a distinct morphology that serves us to improve the biostratigraphic
479 information. Walliser (1964) included in his first Silurian biozonal scheme an *Oz. snajdri*-

480 horizon, which included in the upper part of the *P. siluricus* Zone. Subsequent studies from
481 Gotland and Estonia recognized that “*Oz.*” *snajdri* appeared earlier in the Gorstian (early
482 Ludlow) (Jeppsson *et al.* 1994; Viira and Aldridge, 1998; Jeppsson, 2005). “*Oz.*” *snajdri*
483 becomes extinct in the Cellon Section of the Carnic Alps at the top of the “*Oz.*” *crispa* Zone
484 (Corradini *et al.*, 2015), whereas at the Passo Volajaia area of the Carnic Alps the LAD locates
485 proximate to the base of the lower *O. e. detortus* Zone (Corradini and Corriga 2010; Corradini
486 and Corriga, 2012).

487 The “*Oz.*” *snajdri* Zone aka “*Oz.*” *snajdri* Interval Zone involves the biostratigraphic interval
488 between the LAD of *Polygnathoides siluricus* and the FAD of “*Oz.*” *crispa* (e.g., Corradini
489 and Serpagli, 2009; Cramer *et al.* 2011; Slavík and Carls, 2012; Figure 5). According to
490 Jeppsson *et al.* (2007), the base of the “*Oz.*” *snajdri* Zone is marked by an abrupt major
491 conodont faunal change associated to the improvement of the environmental conditions
492 directly after the Lau event. The lower part of the “*Oz.*” *snajdri* Zone is characterized by the
493 icriodontid *Pedavis latialata* (Walliser) and the upper part by “*Oz.*” *snajdri*. Corradini *et al.*
494 (2015) renamed the “*Oz.*” *snajdri* Interval Zone as *Pedavis latialata* – *Ozarkodina snajdri*
495 Interval Zone for the Cellon Section, considering the two most characteristic species, which
496 co-occur in a short stratigraphic interval.

497 Helfrich (1975) defined the *Spathognathus tillmani* Zone in the lower Wills Creek Formation
498 from the Appalachians based on the occurrence of *S. tillmani* in association with *S. primus*
499 *multidentatus* Helfrich, *Ozarkodina sinuosa* Helfrich, and *O. typica intermedia* Helfrich, and
500 tentatively correlated it with the *Icriodus latialatus* Zone of Walliser (1964). Calner and
501 Jeppsson (2003) regarded *Oz. tillmani* and *S. primus multidentatus* as synonyms of *Oz.*
502 *bohémica* and of *Oz. confluens densidentata* (*S. primus densidentatus* Viira, 1982),
503 respectively. Accordingly, “*Oz.*” *tillmani* would be coeval with *Oz. b. longa*, which ranges

504 from the base of the *Oz. b. longa* Zone and through the *C. murchisoni* Zone, though they are
505 not necessarily conspecific (Calner and Jeppsson, 2003).

506 On the other hand, Viira and Einsnato (2003) noticed that the conodont fauna from the
507 Wenlock–Ludlow transition of Saaremaa is comparable with the faunas described by
508 Helfrich (1975) from the Mifflintown and Wills Creek formations, particularly for the species
509 *Oz. bicornuta* and *Oz. tillmani*. Conodonts from the middle part of the Wills Creek Formation
510 at Pinto, Maryland, merely indicate a Ludlow age (Harris *et al.*, 1994). Successively, the upper
511 part of the Wills Creek Formation and the overlying Tonolway Formation include the index
512 fossil “*Oz.*” *crispa*, which indicates a late Ludlow age (Helfrich, 1975).

513 Following previous studies, “*Oz.*” aff. *snajdri* would correspond to a transitional form of the
514 “*Oz.*” *bohemica* – “*Oz.*” *snajdri* lineage. The record of “*Oz.*” aff. *snajdri* along with *W.*
515 *excavata*, *Ps. beckmanni*, *Ps. bicornis bicornis* suggest a late Homerician – early Gorstian (late
516 Wenlock – early Ludlow) minimum depositional age for the upper debris flows of the
517 Rinconada melange.

518 **TECTONOSTRATIGAPHIC IMPLICATIONS**

519 Silurian deposits of the Precordillera show considerable local and regional variation in
520 thickness and lithology (*e.g.*, Astini and Maretto, 1996; Peralta, 2006, 2007). In the Central
521 Precordillera, the Silurian deposits are widely represented in the La Chilca and Los Espejos
522 Formation of the Tucunuco Group (Cuerda, 1965, 1969), and its lateral equivalent, the
523 Tambolar Formation, which crops out to the south in the Pachaco – San Juan River area
524 (Heim, 1952). Further south and west, Silurian strata are less biostratigraphically constrained.
525 The La Chilca Formation presents in the lower part conodonts from the *Distomodus*
526 *kentuckyensis* Zone (Llandovery, Lehnert *et al.*, 1999), whereas the upper levels include
527 graptolite and acritarch associations assigned to the late Llandovery – early Wenlock

528 (Kerlleñevich and Cuerda, 1986; Pöthe de Baldis, 1987; Albanesi *et al.*, 2006). The Los
529 Espejos Formation paraconformably overlies the La Chilca Formation through a ferruginous-
530 rich basal conglomerate of regional extension. It is succeeded by green sandstone and
531 siltstone packages arranged in a thickening-upward sequence whose thickness ranges
532 between 25 m to 510 m (Cuerda, 1969; Benedetto *et al.*, 1992; Astini and Piovano, 1992).
533 The middle-upper parts of the Los Espejos Formation include in the Jáchal area monospecific
534 associations of *Saetograptus argentinus* (Cuerda) and *Monograptus uncinatus notouninatus*
535 Cuerda, both indicating a Ludlow age (Cuerda, 1969, 1971), and contain conodonts from the
536 *Kokelella v. variabilis* Zone (Gorstian) in the Cerro del Fuerte and the Talacasto areas
537 (Albanesi *et al.*, 2006; Sumrall *et al.*, 2013). García-Muro and Rubinstein (2015) proposed
538 that the middle-upper part of the La Chilca Formation is Wenlock and that the Los Espejos
539 Formation is Wenlock (?), Ludlow to Pridoli, based on palynomorph assemblages and
540 miospore biozones.

541 The iron-rich basal conglomerate, the lithologic contrast and the fossiliferous content suggest
542 the presence of a hiatus between the La Chilca and the Los Espejos formations, involving the
543 late Llandovery – early Wenlock up to the early Ludlow (Peralta, 1993; Astini and Maretto,
544 1996).

545 According to the new conodont fauna from the Rinconada Formation, the hiatus in the
546 Central Precordillera is broadly contemporaneous with destabilization of the eastern
547 Precordilleran platform, accompanying the west-verging thrusting of the Famatinean
548 basement (*e.g.*, Ramos *et al.*, 1996; Mulcahy *et al.*, 2011),

549 **CONCLUSIONS**

550 We analyse here the first Silurian conodont fauna from the Rinconada Formation, proximate
551 to the eastern boundary of the Precordillera. It proceeds from a debris flow from the upper

552 levels of the melange, close to the Neogene disconformity. The conodont association includes
553 the long-ranging species *Dapsilodus obliquicostatus*, *Decoriconus fragilis*, *Pseudooneotodus*
554 *beckmanni*, *P. b. bicornis*, *Wurmiella excavata* along with ozarkodinids of the “Oz.”
555 *bohémica* – “Oz.” *snajdri* lineage. The transitional character of “Oz.” aff. *snajdri* provides a
556 minimum depositional age for the upper debris flows of the Rinconada Formation, probably
557 within the late Homerian – early Gorstian stages. Therefore, the deposition of the upper levels
558 of the Rinconada melange is possibly coeval with the hiatus involved between the La Chilca
559 and Los Espejos formations in the Central Precordillera, and the Silurian deformation in the
560 evolving inner Gondwanan foreland.

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895 **EPIGRAPHS**

896

897 **Figure 1.1.** Geographic location of the Precordillera, Argentina; **2,** Location map of the
898 Rinconada Formation study area, on the eastern flank of the Sierra Chica de Zonda, Eastern
899 Precordillera; **3,** Geographic location of the fossiliferous sample RIN5, Rinconada
900 Formation, San Juan Province.

901

902 **Figure 2.1.** Limestone blocks from the San Juan Formation embedded in debris flows from
903 the upper levels of the Rinconada Formation, proximate to the Neogene unconformity

904 (view to the north); **2**. Iron-rich calcareous level Rin5, which provided the first Silurian
905 conodont fauna from the Rinconada Formation (for location see figure 1); **3**. Debris flow
906 from the upper levels of the Rinconada Formation showing a cleaved matrix with pelitic,
907 siliceous and calcareous intraclasts. Hammer for scale is 33 cm long.

908

909 **Figure 3.** Silurian conodonts from the Rinconada Formation, sample RIN5, Precordillera
910 Argentina. **1–4**, “*Ozarkodina*” aff. *snajdri* (Walliser), **1–2**, Pa element, lateral and oral
911 view, CORD-MP 50913; **3–4**, Pa element, lateral and oral views, see attached coniform
912 element on lateral side, CORD-MP 50914. **5–6**, “*Ozarkodina*” sp., Pa element, CORD-MP
913 50918, lateral and oral views. **7–8, 11**, undetermined “ozarkodinids”, **7**, Pa element,
914 CORD-MP 50941, oral view; **8**, Sa element, CORD-MP 50946, dorsal view; **11**, Pb
915 element, CORD-MP50948, lateral view. **9–10, 17**, *Oulodus* sp., **9**, Sb element, CORD-MP
916 50920, lateral view; **10**, Sb element, CORD-MP 50922, lateral view; **17**, P element, CORD-
917 MP 50943, dorsal view. **12**, *Dapsilodus obliquicostatus* (Branson and Mehl), Sc element,
918 CORD-MP 50815, lateral view. **13, 15, 19**, *Wurmiella excavata* (Branson and Mehl), **13**,
919 Sa element, CORD-MP50921, dorsal (posterior) view; **15**, Pa element, CORD-MP 50919,
920 lateral view; **19**, Sc element, CORD-MP 50923, lateral view. **14**, *Decoriconus fragilis*
921 (Branson and Mehl, 1933), Sb element, CORD-MP 50850, lateral view. **16**,
922 *Pseudooneotodus beckmanni* (Bischoff and Sannemann), P element, CORD-MP 50873,
923 lateral view. **18**, *Pseudooneotodus bicornis bicornis* (Drygant), P element, CORD-MP
924 50870, lateral view. Scale bar: 0.1 mm.

925

926 **Figure 4.** Line-art drawings of selected specimens from sample RIN5. **1–4**, “*Ozarkodina*”
927 aff. *snajdri*, **1–2**, Pa element, lateral and oral view, CORD-MP 50914; **3–4**, Pa element,

928 lateral and oral views, see attached coniform element on lateral side, CORD-MP 50913. **5–**
929 **6**, “*Ozarkodina*” sp., Pa element, CORD-MP 50918, lateral and oral views. **7**, undetermined
930 “ozarkodinid”, Pa element, CORD-MP 50941, oral view. **8–9**, *Wurmiella excavata* (Branson
931 and Mehl), **8**, Pa element, CORD-MP 50919, lateral view; **9**, Sa element, CORD-MP50921,
932 dorsal (posterior) view. **10**, undetermined “ozarkodinid”, Sa element, CORD-MP 50946,
933 dorsal view. Scale bar: 0.1 mm.

934

935 **Figure 5.** Comparison of late Wenlock – Pridoli conodont zones from Estonia (Männik,
936 2014), the Carnic Alps (Corradini *et al.*, 2015), the Prague Synform (Slavík and Carls, 2012;
937 Slavík, 2014), and the Precordillera (this study in grey background), with previous conodont
938 zones of Albanesi *et al.* (2006), Mestre (2009) and Sumrall *et al.* (2013).