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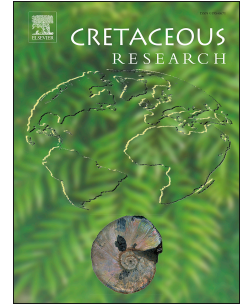
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1 **A revised ammonoid biostratigraphy for the Aptian of NW Africa: Essaouira-Agadir Basin,**
2 **Morocco.**

3

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14

15 **Abstract**

16

17 A revised ammonoid biostratigraphy is presented for the Aptian of NW Africa, Essaouira-Agadir Basin

18 (**EAB**), Morocco, based on detailed analysis of 5 key sections. A number of bio-events are

19 documented and 26 genus and 43 species fully documented, forming the largest published Aptian

20 ammonite collection made from NW Africa. The section at Tiskatine is documented as the type

21 section, and 8 zones and subzones are defined, of which 5 are new. This work allows correlation of

22 the Aptian of the **EAB** to the Standard Mediterranean Ammonite Scale (**SMAS**).

23 Two main hiatuses are identified at the scale of the basin scale: a major one that includes most of

24 the lower Aptian and the base of the upper Aptian and a second one encompass the top of the

25 upper Aptian and the base of the lower Albian. The ammonite fauna displays a clear Tethyan

26 palaeobiogeographic character affected by a fairly high degree of endemism at the genus and

27 species level. The new genus and species *Elsaella tiskatinensis* is introduced.

28

29 **Keywords:** Ammonite, Biostratigraphy, Aptian, Essaouira-Agadir Basin, Morocco

30 **1. Introduction**

31

32 Aptian to lowermost Albian ammonite biostratigraphy across the northwest African Atlantic
33 margin remains poorly documented compared to counterparts of the Mediterranean Tethys; such as
34 in Spain (Moreno Bedmar et al., 2008, 2009, 2010, 2012a, 2014), France (Delanoy, 1995a; Cecca et
35 al., 1999; Ropolo et al., 1999; Kennedy et al., 2000; Ropolo et al., 2000a-c; Dauphin, 2002; Dutour,
36 2005; Ropolo et al. 2006; Ropolo et al., 2008a-b; Frau et al., 2015), Tunisia (Lehmann et al., 2009;
37 Chihaoui et al., 2010; Latil, 2011), and Iran (Raisossadat, 2004, 2006; Bulot in Vincent et al., 2010;
38 Seyed-Emami and Wilmsen, 2016); or on the opposite side of the Central Atlantic Margin (CAM), e.g.
39 Mexico (Barragán-Manzo and Méndez-Franco, 2005; Moreno Bedmar et al., 2012b, 2013, 2015,
40 Barragán et al., 2016).

41 The Aptian Stage contains some globally significant events, recording a time of high sea-level
42 during the mid-Cretaceous greenhouse period (Larson and Erba, 1999; Leckie et al., 2002). It also
43 records some of the most wide-spread and best studied oceanic anoxic events (Schlager and
44 Jenkyns, 1976; Scholle and Arthur, 1980; Arthur et al., 1990; Bralower et. al, 1994; Weissert et al.,
45 1998; Föllmi et al., 2006; Föllmi, 2012).

46 In this study we present a new high-resolution dataset from NW Morocco, discussed within
47 a broader regional/global context and with special reference to the standard ammonite zonal
48 scheme of the Mediterranean Tethys (Reboulet et al., 2011, 2014). In Morocco, the Aptian
49 successions of the west-central part of the Essaouira-Agadir Basin (**EAB**) is part of one of the most
50 complete and best-constrained successions of the Lower Cretaceous in NW Africa. The Aptian
51 succession is characterized by ammonite-rich, shallow-marine (Bouzerrou Formation) to shelf
52 deposits (Tamzergout and Oued Tidzi formations), that crop out in a corridor along the Moroccan
53 coast between the cities of Agadir and Essaouira (Fig. 1). The aim of the present contribution is to
54 improve the biostratigraphic frame of the Aptian and lowermost Albian of the **EAB** based on the
55 detailed analysis of the bed by bed distribution of ammonoids at five key sections (Fig. 1).

56 Our results document the specific character of the ammonoid fauna of the **EAB** and discuss
57 its similarities and differences with equivalent ammonite faunas of Mediterranean-Caucasian
58 Subrealm of the Tethyan Realm. The results of this study develop a strong reference framework for
59 the Aptian to earliest Albian for the northwest African Atlantic margin (figure 12).

60

61 **2. Regional Setting and Stratigraphy**

62

63 Lower Cretaceous strata crops out over an area 100 km wide and 150 km long, adjacent to
64 the Moroccan Atlantic coast; the only continuous succession of Mesozoic fill of the Central Atlantic
65 Margin (CAM) (Fig. 1). Following Permo-Triassic rifting that separated Africa and North America (Le
66 Roy and Piqué, 2001), extensive Jurassic carbonate platforms developed all along the NW Africa
67 (Jansa and Wiedmann, 1982). Later deposition shared this inherited regional physiography, but also
68 record distinct sub-basins with variable subsidence, evolving throughout the Early Cretaceous (Rey
69 et al., 1988; Davison, 2005; Wenke et al., 2011).

70 During this time, the **EAB**, the focus of this study, was limited to the north by the Meseta
71 and the Jebilet, to the south by the Souss Basin and to the east by the Massif Ancien de Marrakech.
72 The general physiography formed a large gulf-like embayment opening out to the west into the
73 Atlantic Ocean (Behrens et al., 1978). Present day exposure of the Mesozoic basin fill and adjacent
74 basement terrains reflects the Cretaceous to Cenozoic inversion and uplift linked to the
75 Atlasic/Alpine orogeny (Laville et al., 2004) and associated salt tectonics (Tari and Jabour, 2013).

76 This study focuses on the outcrops in the west-central part of the **EAB**, located between the
77 prominent Cretaceous Essaouira and Chichaoua escarpment to the north and the broad Cap Ghir
78 anticline to the south (Fig. 1). Strata are mainly flat lying, only exhibiting folding with steep flanks
79 close to fault zones and associated with salt diapirs (Hafid et al., 2000, 2006). The maximum
80 thickness of the entire Cretaceous post-rift succession is estimated to be 1300 m (Behrens et al.,
81 1982) (Zem Zem section). The Lower Cretaceous reaches approximately 800 m in thickness but

82 varies dramatically, thinning toward salt diapirs, suggesting Cretaceous syn-sedimentary activity
83 (Hafid et al., 2000; Le Roy and Pique, 2001; Zühlke et al., 2004).

84 The Early Cretaceous represents a time of major change from the predominately carbonate-
85 dominated system in the Middle- to Late Jurassic to a mixed carbonate-siliciclastic or purely
86 siliciclastic systems in the Cretaceous, following drowning of the Jurassic carbonate platform.
87 Cretaceous deposits in the **EAB** are dominated by shelfal mudstone successions with discrete
88 intervals of coarse clastic sediment delivery in the late Valanginian, late early Hauterivian and late
89 Barremian to earliest Aptian (Ambroggi, 1963; Behrens et al., 1978).

90 During the Aptian the paleogeography of the **EAB** was dominated by an open marine shelf.
91 Arid climatic conditions are indicated by low nannofossil productivity (Herrle, 2002; Peybernes et al.,
92 2013) and clay mineral composition (Daoudi and Deconinck, 1994). The **EAB** is thought to have been
93 located away from the main upwelling zone, which existed during Aptian and early Albian times
94 around the Mazagan Plateau to the north (Herrle, 2002). A cooling climatic trend has been
95 recognized from the late Aptian – early Albian, evidenced by southward migration of high-
96 latitude/boreal nannofossil into mid- and low-latitudes (Jeremiah, 1996, 2001; Herrle and
97 Mutterlose, 2003; Rueckenheim et al., 2006). This trend was also observed in the **EAB** by Peybernes
98 et al. (2013).

99 Overall the Aptian to Albian transition is a time of global eustatic sea-level rise (Hardenbol et
100 al., 1998; Haq, 2014). In the **EAB** shallow-marine conditions were widespread during the early
101 Aptian, with a transgression recognised in the late Aptian, the establishment of outer shelf
102 conditions, and reoccurrence of the Atlas Gulf (Behrens et al., 1978).

103 Most of the formation names for the Lower Cretaceous were introduced by Duffaud et al.
104 (1966), later revised by Rey et al. (1986a and b, 1988). The reference sections are mainly located
105 along an E-W trending transect in the northern part of the **EAB** (Essaouira to Imi'n'Tanoute). There
106 are no stratigraphic units at group level defined in the **EAB**. Previous sedimentological and
107 stratigraphic work on the Lower Cretaceous in the areas was carried out by Ambroggi (1963),

108 Duffaud et al. (1966), Wiedmann et al. (1978, 1982), Adams et al. (1980), Behrens et al. (1982) and
109 Rey et al. (1986a and b, 1988). More recent studies have mainly focused on the ammonite
110 biostratigraphy of the Berriasian to Hauterivian interval (Ettachfini, 1991, 2004; Wippich, 2001,
111 2003) and the integrated stratigraphy of the Barremian/Aptian interval (Witam et al., 1993; Witam,
112 1998; Nouidar and Chellai, 2001, 2002; Company et al., 2008; Peybernes et al., 2013).

113

114 **2.1 Barremian to Albian lithostratigraphic framework**

115

116 The first attempt to subdivide the lithostratigraphic succession of the **EAB** was made by
117 Duffaud et al. (1966). For the Barremian to Albian interval, those authors introduced five
118 lithostratigraphic units: the "Calcaires lumachéliques de Taboulaourt", the "Grès et marnes rouges
119 du Bou Zergoun", the "Marno-calcaires de Tamzergout", the "Grès marneux du Lemgo" and the
120 "Marnes de l'Oued Tidsi". This synoptic lithostratigraphic chart was introduced without formal
121 description, a definition of boundaries or designation of reference sections.

122 Subsequent works by Rey et al. (1986a and b, 1988), Andreu (1989), Witam et al. (1993) and
123 Witam (1998) led to refining the litho- and biostratigraphic framework of the **EAB**, but little
124 attention was paid to the formal description and definition of the formations. Despite Witam's
125 (1998) attempt to propose reference sections for the lithostratigraphic units introduced by Duffaud
126 et al. (1966), a unified lithostratigraphic nomenclature for the **EAB** is still lacking. This drastically
127 limits the value of the lithostratigraphic units for the correlation of the Barremian to Albian strata
128 and also reflects the limited published work on regional stratigraphic relationships. Revising the
129 lithostratigraphic framework of the **EAB** is beyond the scope of the present contribution and in this
130 study we have utilised the existing scheme of Duffaud et al. (1966) that best applies to our
131 observations in the west-central part of the basin (Fig. 2).

132

133 *2.1.1 Bouzergoun Formation*

134

135 The Bouzergoun Fm., composed of sandstones and red mudstones, was introduced by
136 Duffaud et al. (1966). Rey et al. (1986a, 1988), describes them as margino-littoral deposits made of
137 sands, varicoloured clays, dolomites and bioclastic-rich limestones with large cross-stratification;
138 and topped by a major disconformity.

139 There is no agreement in the literature about the age of the Bouzergoun Fm. Rey et al.
140 (1986a) proposed a late Barremian age, based on the occurrence of early Barremian ammonites in
141 the underlying Taboulouart Fm. and early Aptian ammonites in the overlying Tamzergout Fm. This
142 was amended by Nouidar and Chellaï (2001) who assigned the upper part of the Bouzergoun Fm.
143 (red beds) to the lower Aptian, based on the occurrence of *Salpingoporella? dinarica* Radoičić.
144 However, it is unclear if the occurrence is derived from new data obtained by the authors or from a
145 misinterpretation of Canérot et al. (1986), who never reported these dasyclad algae from the **EAB**. In
146 any case, the occurrence of *Salpingoporella? dinarica* does not allow discrimination between the
147 Barremian and Aptian stages.

148 Analysis of outcrops for this new study identify the Bouzergoun Fm. as being composed of a
149 thick succession of shelfal muds with minor sandstones at the base, truncated by coarsening and
150 thickening-upward shallow-marine and deltaic sand-rich deposits and fluvial deposits. In places,
151 erosional channel and valley features are in-filled by coarse clastic material and greenish to red-
152 coloured mudstones with interbedded sandstones. The uppermost part of the formation is made of
153 sandstones, sandy limestones and mudstone interbeds bearing marine fauna and often oyster-rich
154 beds to the top (Fig. 2). The unit reaches a maximum thickness of 84 metres at Assaka.

155 Ammonites collected from Assaka identify a condensed horizon that marks the boundary
156 between the Taboulouart and the Bouzergoun formations, of early late Barremian age (early to
157 middle part of the *Gerhardtia sartousiana* Zone) (see also discussion in Company et al., 2008). In
158 most sections studied the occurrence of *Procheloniceras dechauxi* (Kilan and Reboul, 1915) firmly
159 establish an earliest Aptian age (see discussion below) for the uppermost beds of the Bouzergoun

160 Fm. (see discussion below). At Tiskatine, the top of the formation is marked by two beds of siltstones
161 that contain a rich early late Aptian ammonite fauna (lower part of the *Colombiceras tobleri* Zone)
162 (see discussion below). This indicates that the top of the Bouzergoun Fm. is diachronous and ranges
163 in age from earliest to early late Aptian (base of the *Deshayesites forbesi* to lower part of the
164 *Colombiceras tobleri* zones).

165

166 2.1.2 Tamzergout Fm.

167

168 The Tamzergout Fm. was defined by Duffaud et al. (1966) as a succession of marls and
169 limestones. The vertical and lateral extension of this unit has been variously interpreted by
170 subsequent authors (see discussion in Witam, 1998, p. 153-154).

171 The formation is mainly made up of fossiliferous alternating blue-grey marls and grey
172 limestones that correspond to shallow-marine to shelfal deposits (Fig. 2). Rey et al. (1986a)
173 restricted its age to the early Aptian (= Bedoulian in regional French stratigraphy) and limited its
174 occurrence to the west and central part of the **EAB**. Rey et al. (1986a) also introduced the Tadhart
175 Fm. of late Aptian age (= Gargasian in regional French stratigraphy), sitting between the Tamzergout
176 and subsequent Oued Tidzi Fm. In this study we propose a different age range for the Tamzergout
177 Formation (see below), from the early to latest Aptian. We therefore consider the Tadhart Fm. a
178 time-equivalent unit for parts of the upper Tamzergout Fm. that is restricted to the eastern and
179 more proximal parts of the basin. In the western and central part of the basin it cannot be
180 discriminated from the Tamzergout Fm. In the studied area, the top of the Tamzergout Fm. is
181 marked by a regional unconformity that was previously reported from Agadir (discontinuity **D4** of
182 Peybernes et al. 2013). The formation reaches a maximum thickness of 33 metres at Tiskatine.

183 In most sections studied, the lower part of the formation contains the base of the
184 *Deshayesites forbesi* Zone (lowermost Aptian). *Procheloniceras dechauxi* is abundant, but first
185 appears in the underlying Bouzergoun Fm. At Tiskatine, the first limestone bed of the Tamzergout

186 Fm. is marked by the First Occurrence (**FO**) of "*Epicheloniceras*" *marocanus* (Roch, 1930) that
187 indicates the base of the upper part of the *Colombiceras tobleri* Zone (see discussion below). The
188 assigned ages further highlight the diachronicity of the top Bouzergout and base Tamzergout
189 Formation. The upper part of the formation is usually rich in small pyritic ammonites that indicate a
190 latest Aptian age (*Elsaella tiskatinensis* Zone, see definition and discussion below).

191

192 2.1.4 Lemgo Formation

193

194 The Lemgo Fm. was introduced by Duffaud et al. (1966) and subsequently reinterpreted by
195 Rey et al. (1986b) to include a complex of green marls, yellow sandy marls and sandy dolomites at
196 the top. It is named after the Jbel Lemgo ridge, to the east of Imi'n'Tanoute.

197 Based on our field observations and microfacies analysis in the Mramer section, the Lemgo
198 Fm. is composed of argillaceous sandstones and bioclastic-rich sandy limestones. Regional
199 correlations suggest that the Lemgo Fm. represents the proximal equivalent of the upper
200 Tamzergout Formation and is not present in the western central part of the basin (see discussion
201 below).

202 Rey et al. (1988) assigned the formation to the uppermost Aptian (= Clansayesian in regional
203 French stratigraphy) based on the occurrence of ammonite assemblages that characterized the
204 *Nolaniceras nolani* and *Hypacanthoplites jacobi* zones. Material collected in this study of what is
205 exposed of the Lemgo Fm. at the Mramer section indicates a latest Aptian to earliest Albian age
206 (*Acanthohoplites ashiltaensis* to *Mellegueiceras chihaouiae* zones, see discussion below).

207

208 2.1.4 Oued Tidzi Formation

209

210 Described by Duffaud et al. (1966) as marls, it was redefined by Rey et al. (1988) as a
211 complex of green marls bearing small pyritic ammonites, intercalated with marly limestones and
212 sandy dolomites.

213 The unit is easily recognized in the west-central part of the **EAB** where it reaches a maximum
214 thickness of 340 m and forms extensive recessive slopes in the landscape. Our field observations and
215 microfacies analysis, focused on the lower part of this formation, show a clear change from blue
216 marls and limestones of the Tamzergout Fm. to a complex of green marls with minor bioclastic-rich
217 sandstone interbeds (Fig. 2). The early Albian age of the basal sandstones is established by the
218 occurrence of the diagnostic ammonite genus *Douvilleiceras*.

219

220 **2.2 Previous Work on Aptian biostratigraphy**

221

222 Pioneering work reporting and describing ammonites in the **EAB** was undertaken by Kilian
223 and Gentil (1906), Roch (1930) and Ambroggi (1963). The work of Ambroggi (1963) has to be
224 highlighted, as it established the first regional biostratigraphic framework for the Lower Cretaceous.
225 The ammonite biostratigraphy of the Berriasian - Barremian interval was subsequently refined by
226 Ettachfani (1991, 2004), Wippich (2001, 2003) and Company et al. (2008). Yet, despite the good
227 accessibility of sections and abundance of ammonites, no recent studies have focused on the
228 detailed ammonite palaeontology and biostratigraphy of the Aptian – Albian interval.

229 Following the work of Roch (1930) and Ambroggi (1963), extensive faunal lists of Aptian and
230 Albian ammonites were published by Bergner et al. (1982), Rey et al. (1986a and b, 1988), Witam et
231 al. (1993) and Witam (1998). Unfortunately, the precise stratigraphic position of the material within
232 the formations was not documented and only a very limited number of specimens were illustrated in
233 the literature. As a consequence, the available previously published data sets do not fulfil the
234 standards of modern biostratigraphic studies and it is difficult to reinterpret them in the light of our
235 own results.

236 Most recently a detailed biostratigraphic framework based on bed by bed collections from a
237 transect along the Agadir segment of the **EAB** was published by Peybernes et al. (2013). A fairly high
238 degree of endemism was suggested by the introduction of many new species and genus names that
239 are unfortunately not formally described or illustrated. The authors chose not to introduce new
240 biostratigraphic units, but pointed out the necessity for a local ammonite zonal scheme. The
241 published zonation is an attempt to apply the upper lower Aptian to lower Albian ammonite zonal
242 scheme of northern Tethys (Reboulet et al., 2011) to the **EAB**. This interval is described as being
243 strongly affected by condensation and temporal hiatuses. According to these authors, the co-
244 occurrence of *Chelonicerias sp.* and *Deshayesites sp.* indicates an early Aptian age for the base of the
245 Tamzergout Fm. (tentatively assigned the *Deshayesites deshayesi* Zone). Condensation and merging
246 of unconformities across the lower/upper Aptian boundary is supported by the mixing of *Dufrenoyia*
247 *furcata* and *Epicheloniceras martini* zones assemblages. The upper Aptian *Epicheloniceras martini*,
248 *Parahoplites melchioris*, *Acanthohoplites nolani* and *Hypacanthoplites jacobi*, zones are identified on
249 the basis of characteristic ammonites assemblages, even so it is highlighted that the extension of the
250 *P. melchioris* Zone is uncertain, as the index fossil is absent and indicative taxa are scarce. The lower
251 Albian *Leymeriella tardefurcata* Zone and *Douvilleicerias mammillatum* Superzone are recognized
252 throughout most of the basin even though the index species are absent. It should be noted that a
253 bed by bed distribution of ammonites is only documented in the Addar section and the authors do
254 not provide photographic plates to substantiate the ammonite systematics used in their
255 contribution.

256

257 **3. Studied Sections**

258

259 The present publication is focused on five Aptian-Albian sections (Fig. 1). Locations were
260 chosen to reinvestigate previously studied sections for reference and further to add new sections for
261 better spatial constraints and coverage of the western, central and the northern part of the basin.

262

263 **3.1. Tiskatine** (Fig. 3) - Lat.: 30.821463° Long.: -9.702555° (Tiskatine 1) and Lat.: 30.810477° Long.: -
264 9.739966° (Tiskatine 2).

265

266 The lower part of the succession (beds **TK 159 to 206**, Fig. 3) is best exposed 6.5 Km to the
267 east of the village of Assaka (Tiskatine 1), northwest of Adrar (mountain) Tiskatine. The upper beds
268 (**TK 206 to 249**) are better exposed 3 Km to the east of Assaka (Tiskatine 2). The two sections were
269 correlated using the marker bed **TK 206**. Section Tiskatine 1 starts in the river bed south of the road
270 from Assaka to Tazzougart and continues to the north. At Tiskatine 2 the beds crop out on both sides
271 of the road. Tiskatine 1 was previously studied by Roch (1930) and Ambroggi (1963).

272

273 3.1 m of the upper part of the Bouzergoun Fm. are exposed at Tiskatine. The first bed (**TK**
274 **157**) is a well-consolidated oyster-rich, ammonite-barren, sandy limestone topped by an iron-rich
275 crust. It is followed by yellow sandstones interbedded with yellow to green mudrock partings (beds
276 **TK 158-162**). The sandstones are fine-grained and show low angle cross-bedding and laterally
277 extensive undulating surfaces. They are often topped by iron-enriched crusts, yielding phosphatic,
278 glauconitic pebbles and fossils. The fossil content comprises ammonites, belemnites, echinoids,
279 gastropods, and rare solitary coral fragments. The top surface of bed **TK 161** is marked by a
280 belemnite accumulation that suggests transport and winnowing by currents (type 4 condensation
281 accumulates of Doyle and Macdonald, 1993). The deposits represent a deepening-upward shallow
282 marine succession. Wave-influenced shoreface sedimentation dominates at the base (bed **TK 157**)
283 with a subsequent transition into lower shoreface to offshore sedimentation (bed **TK 158-162**). The
284 contact with the overlying Tamzergout Fm. is marked by well-developed marls in bed **TK 163**. The
285 contact is abrupt and marks the change to shelfal conditions below storm wave base.

286

287 The alternating blue-grey marls and grey limestones of the Tamzergout Formation develop
288 from bed **TK 163** to bed **TK 247**. The marls are laminated, forming decimetre to metre thick beds.
289 Limestones are 10 to 20 centimetres thick and appear massive. Contact between alternating beds
290 are mostly sharp and planar. In the middle part of the succession some limestones beds have detrital
291 component (**TK 180, 182, 192** and **194**) and two prominent fine-grained sandstone beds (**TK 184** and
292 **TK 186**) are recognised. The sandstones are topped by ferruginous crusts. From bed **TK 197** to bed
293 **TK 246** the succession is characterized by a fairly monotonous alternation of marls and limestones.
294 **TK 206** is a prominent extremely fossiliferous marker bed, with a well-developed iron-crust on the
295 top. Above, the marls are rich in pyritic nodules and ammonites, and some minor detrital input was
296 observed toward the top of the Tamzergout Fm. that ends at a distinct glauconitic horizon (bed **TK**
297 **247**) rich in phosphate nodules. The total thickness of the Tamzergout Fm. is 33.15 metres.
298 Sedimentation of the Tamzergout Fm. is mainly through suspension fall out. Sandstone beds **TK 184**
299 and **TK 186** are interpreted as gravity flow deposits.

300 The base of the Oued Tidzi Fm. is marked by a fossil-rich sandstone bed **TK 248** that contains
301 reworked phosphatic pebbles and glauconitic fossils (including ammonites). It is capped by a well-
302 developed iron crust directly overlain by distinctive green to yellow marls of the Oued Tidzi Fm. Bed
303 **TK 248** is interpreted as a shelf gravity flow.

304 Ammonite abundance and diversity vary throughout the succession. This discontinuous
305 palaeontological record may partially reflect collection failure, despite an intensive search of the
306 barren intervals. Based on our present knowledge, the following sequence of bio-events is
307 recognised as regionally significant following comparison with the other sections studied:

308

309 - Bio-event 1: sudden mass occurrence of *Procheloniceras* in bed **TK 159**, followed by the first
310 observed occurrence of *Deshayesites* in bed **TK 160**;

311 - Bio-event 2: peak of diversity in bed **TK 161a** dominated by *Colombiceras* and
312 *Epicheloniceras*;

- 313 - Bio-event 3: first apparition datum of the endemic species "*Epicheloniceras*" *marocanus* at
314 the top of bed **TK 162**, followed by its acme in bed **TK 166**;
- 315 - Bio-event 4: lowest occurrence of *Acanthohoplites* in bed **TK 184**;
- 316 - Bio-event 5: peak of abundance of *Pseudoaustraliceras* in beds **TK 192** and **TK 194**;
- 317 - Bio-event 6: sudden mass occurrence of *Elsaella* gen. nov. in bed **TK 206**;
- 318 - Bio-event 7: radiation of the Acanthohoplitidae (*Nodosohoplites*, *Protacanthoplites* and
319 "*Hypacanthoplites*") from bed **TK 206** to bed **TK 221**;
- 320 - Bio-event 8: lowest occurrence of *Douvilleiceras* in bed **TK 248**.

321

322 Tiskatine is the only studied section where the complete succession of events was
323 recognized and documented. It has, therefore, been selected as the reference section for the
324 ammonite biostratigraphy of the west central part of the **EAB**.

325

326 **3.2. Tamanar** (Fig. 4) - Lat.: 31.057403° Long.: -9.601503°

327

328 This outcrop is 10 Km northeast of Tamanar and the section is also known as Ida Ou Shak
329 named after the nearby village. It can be reached via an unmade track that leaves National road 1
330 approximately 8 Km north of Tamanar. This section has not been previously described in literature.

331 Fine-grained laminated sandstones with interbedded calcareous mudstones make up the
332 lower 8 metres (beds **TM 49** to **58**) of the studied section and comprise the uppermost part of the
333 Bouzergoun Fm. Oysters and other large bivalves are abundant throughout. The lower part of the
334 logged section is affected by soft sediment deformation. The first occurrence of ammonites is
335 recognised on top of the slumped interval. The deposits of the Bouzergoun Fm. here are interpreted
336 as lower shoreface to offshore shelf transition.

337

338 The base of the Tamzergout formation is marked by the change to limestones and
interbedded marls, both containing abundant open-marine fauna (e.g. ammonites and belemnites).

339 The formation encompasses beds **TM 59** to **TM 80**. Compared to the interval exposed at Tiskatine,
340 the limestones here have a stronger siliciclastic detrital component. The succession is fairly
341 monotonous, but a few beds have notable features. From bottom to top, these are:

342

343 - Bed **TM 69** contains very abundant ammonites (dominantly *Procheloniceras dechauxi*)
344 associated with a high ferruginous content;

345 - Bed **TM 70a** is a slumped complex marked the only occurrence of upper lower Aptian
346 ammonites (*Cheloniceras*) identified in this study;

347 - Bed **TM 71** contains a phosphatic, glauconitic and ferruginous fossil assemblage rich in
348 ammonites, belemnites, bivalves, and brachiopods.

349 - Bed **TM 73** contains belemnites and bivalves and is marked by a well-developed and
350 prominent iron crust.

351 - Bed **TM 76** is rich in ammonites and marks an increase in diversity of the ammonite fauna

352 - Bed **TM 80** yields abundant pyritic ammonites.

353

354 The sandstone of bed 81 is interpreted to define the base of the Oued Tidzi Formation,
355 displaying a similar depositional environment to Tiskatine.

356 The Tamzergout Formation has a total thickness of 19.20 metres. At this location it is
357 interpreted to be mixed shelfal pelagic marls, with changes in siliciclastic sediment influx transported
358 by wave and current supported gravity flows, forming the sandy limestones.

359 Compared to the succession of bio-events recognized at Tiskatine, several points should be
360 outlined:

361

362 - Bio-event 1 is marked by a less sudden appearance of *Procheloniceras* in bed **TM 55**;

363 - Bio-event 2 is not well expressed due to poor outcrop conditions but merely lies at the top of
364 bed **TM 71**;

- 365 - Bio-event 3 is marked by the spot occurrence of "*Epicheloniceras*" *marocanus* in bed **TM 72**;
366 - Bio-event 4 and 5 are not recorded;
367 - Bio-event 6 and 7 are identified in beds **TM 76** to **TM 80** with a slightly different expression
368 due to condensation combine with temporal hiatus.

369

370 **3.3. Zem Zem** (Fig 5) - Lat.: 31.241846° Long.: -9.372052°

371

372 Located about 17 Km to the south-southeast of Meskala, this outcrop can be reached via an
373 unmade road leaving the main road between Bizdad and Ait Daoud, approximately 2 Km south-
374 southeast of Bizdad. Nearby sections have been studied by Roch (1930), Butt (1982), Rey et al.
375 (1986a, 1988) and Witam (1998).

376 The Bouzergoun Fm. reaches a thickness of 12 metres at Zem Zem (up to bed **ZZ 60**). The
377 upper part is composed of sandy rudstones, sandstones and minor wackestones with marls
378 interbedded. Fauna is dominated by brachiopods, oysters, ammonites and echinoderms, pointing
379 towards open-marine conditions. Horizon **ZZ 59** is a prominent ferruginous surface just below the
380 topmost bed of the formation, containing abundant fossils with phosphatic preservation. Deposition
381 overall took place in a shoreface to carbonate-rich subtidal environment with higher energy index,
382 on a shallow shelf.

383 The Tamzergout Fm. is interpreted to extend from bed **ZZ 61** to **ZZ 68** and is dominated by
384 marls with two prominent limestone beds (**ZZ 61** and **ZZ 65**) and a sandstone bed topped by an iron
385 crust (**ZZ 67**). The thickness of the Tamzergout Fm. is approximately 20 metres, and the presence of
386 laterally extensive marls with limestone interbeds containing open marine fauna and reduced
387 siliciclastic input suggests a low-energy shelf environment, similar to that at Tiskatine.

388 The base of the Oued Tidzi Fm. is defined as a prominent sandstone horizon, and continues
389 to the top of the measured section, with a minimum thickness of 10 metres.

390 At Zem Zem, the palaeontological record shows that bio-event 1 (mass occurrence of
391 *Procheloniceras*) is well expressed and almost directly overlain by the sudden appearance of *Elsaella*
392 gen. nov., followed by the radiation of the Acanthohoplitidae (bio-events 6 and 7). Bio-event 2 is
393 suspected in the condensed horizon at the top of the Bouzergoun Fm. (spot occurrence of
394 *Epicheloniceras* with reworked *Deshayesites* at the top of bed **ZZ 60**). Bio-events 3, 4 and 5 are not
395 recorded and outline the existence of a major temporal hiatus at the base of the Tamzergout Fm.

396

397 **3.4. Oued Tlit** (Fig. 6) - Lat.: 31.194306° Long.: -9.665616°

398

399 The section is located 4 Km southeast of Smimou, on the northern flank of the Djebel
400 Amsittene anticline, north of the road between Smimou and Imi'n'Tlit. This locality has previously
401 been studied by Roch (1930), Rey et al. (1986a, 1988) and Witam (1998).

402 The top of the Bouzergoun Formation comprises the lowermost 2.10 metres. It is composed
403 of fine to medium-grained non-amalgamated sandstones interbedded with mudstones. The
404 sandstones exhibit hummocky cross-stratification and are often affected by in-situ soft-sediment
405 deformation. The upper part is formed by a very prominent set of oyster rudstones. The depositional
406 environment is interpreted as shoreface to subtidal.

407 Ammonites occur immediately above the set of oyster rudstones and the remaining 11.70 m are
408 interpreted to as part of the Tamzergout Formation. It comprises alternating marls and micritic
409 limestones. A well-expressed hardground surface marked by iron crust and glauconite, associated
410 with belemnite accumulation and phosphatic ammonites, occurs at the top of bed **OT 11**. The upper
411 part of the Tamzergout Fm. has not been investigated in this study.

412 The succession of bio-event 1 (sudden appearance of *Procheloniceras*) and bio-event 2 (peak
413 abundance of *Epicheloniceras* and *Colombiceras*) are recognized in the lower part of the Tamzergout
414 Fm. Our findings also question the identification of the *Dufrenoyia* and *Epicheloniceras* illustrated by
415 Witam (1998) from this section (see discussion below).

416

417 **3.5. Mramer** (Fig. 7) - Lat.: 31.657033° Long.: -9.164205° (Mramer 1) and Lat.: 31.666118° Long.: -

418 9.153251° (Mramer 2).

419

420 Mramer represents the northernmost section studied in the **EAB**. It is located approximately

421 60 Km to the ENE of Essaouira and 12 Km NE of Tafetachte. The Mramer section is a composite

422 section, both locations cropping out in a river bed. Mramer 1 is located about 1.5 Km to the

423 northeast of the market in Mramer and location Mramer 2 is at the base of a steep slope south of

424 the market. This outcrop was mentioned by Roch (1930) but not reported in detail.

425 The upper 3.50 metres of the Bouzergoun Fm. are exposed at this locality. Strata are

426 dominated by sandstones and siltstones. The fauna is rich in oyster fragments, and further includes

427 scarce ammonites; towards the top echinoderm fragments are common. The medium to fine-

428 grained sandstones exhibit in-situ soft sediment deformation structures and common bioturbation,

429 they are dominated by low-angle cross lamination, interpreted to have been deposited under

430 shoreface conditions.

431 The following interval (beds **MR 9a** to **9f**) comprises part of the Tamzergout Fm., here

432 composed of sandy limestones and marls interbedded. The limestones contain belemnites, bivalves,

433 and echinoderms and exhibit prominent iron crusts at the tops. The exposed part of the Tamzergout

434 Fm. records a return to shelfal, open water, and lower-energy depositional environments. This

435 interval is capped by a zone of poor exposure that is 17.5m thick and likely composed of limestones

436 and interbedded marls with increasing abundance of ammonites.

437 It is overlain by the Lemgo Fm., the proximal equivalent to the upper Tamzergout Fm. The

438 base of the Lemgo Fm. is not exposed in the study area. It is composed of yellow argillaceous

439 sandstones and sandy limestones obtaining a minimum thickness of 4.50 metres. Fossil-rich sandy

440 limestones at the base (beds **MR 9x** to **9z**) are laterally continuous. Beds **MR 10** to **MR 17** comprise

441 very fine to fine-grained sandstones interbedded with floatstones. Sandstones exhibit large-scale,

442 higher energy dune bed forms and often rework the interbedded limestones. Limestones have well-
443 developed iron crusts and are rich in fossils containing brachiopods, ammonites, gastropods, and
444 belemnites. These fossils often occur in phosphatic and glauconitic preservation, pointing to a
445 reoccurrence of anoxia.

446 This mixed siliciclastic-carbonate succession is interpreted to represent shelfal carbonates
447 being reworked by sandstones deposited during intermittent, wave-influenced sedimentation.
448 Abundance of ammonites and cosmopolitan taxa imply an open –marine connection but
449 reoccurrence of anoxia and absence of indicative, open-marine fauna is pointing towards a periodic
450 development of restricted environment. Overall, this represents a shallowing-up succession with a
451 higher energy index than other time-equivalent sections studied in the sections of the central and
452 western part of the basin. The Lemgo Fm. is overlain by marls and minor sandy limestones of the
453 Oued Tidzi Fm. that starts at bed **MR 20** and marks the return into lower-energy shelfal
454 sedimentation.

455 At Mramer, the Lemgo Fm. records the local expression of bio-events 6 to 8. The ammonite
456 succession is characterized by the radiation of the Acanthohoplitidae from bed **MR 10m** to **MR 15**,
457 followed the lowest occurrence of *Douvilleiceras* in bed **MR18**. Noteworthy is the occurrence of a
458 well preserved and abundant fauna of *Mellegueiceras*, a genus that was so far only known from
459 Central Tunisia (Latil, 2011).

460

461 **4. Systematic Palaeontology**

462

463 More than 1000 ammonite specimens were collected during two field sessions in 2015 and
464 2016. To our knowledge, the material collected represents the largest bed by bed collection made
465 from the Bouzergoun and Tamzergout formations of the central and northern part of the **EAB**. The
466 preservation of the material is variable and includes internal calcareous, phosphatic and pyritic

467 moulds. Abundance varies considerably throughout the studied sections and high diversity is
468 observed in specific horizons, some of which can be correlated at the scale of the **EAB**.

469 As already outlined in the faunal lists given by our predecessors (Roch, 1930; Ambroggi,
470 1963; Rey et al., 1986a), the ammonite assemblages are largely dominated by the Douvilleiceratidae
471 Parona and Bonarelli, 1897 and Acanthohoplitidae Stoyanow, 1949. The Deshayesitidae Stoyanow,
472 1949, Desmoceratidae Zittel, 1895 and Ancyloceratidae Gill, 1871 are minor elements of the faunas.
473 Aconeceratidae Spath, 1923, Phylloceratidae Zittel, 1884 and Tetragonitidae Hyatt, 1900 are even
474 rarer.

475 Despite an extensive list of published work, the systematics of the Aptian Ammonoidea is
476 still at a very preliminary stage due to the lack of modern taxonomic revision taking in consideration
477 intraspecific variation and sexual dimorphism. This is especially true for the Douvilleiceratidae and
478 Acanthohoplitidae, for which a plethora of typological species was introduced over the years (see
479 lists in Klein and Bogdanova, 2013). Moreover, there is hardly any agreement among authors
480 regarding the limits and content of *Colombiceras* Spath, 1923, *Acanthohoplites* Sinzow, 1908;
481 *Protacanthoplites* Tovbina, 1970; *Nolaniceras* Casey, 1961; *Hypacanthoplites* Spath, 1923;
482 *Procheloniceras* Spath, 1923; *Diadochoceras* Hyatt, 1900; *Nodosohoplites* Egoian, 1965 and
483 *Epicheloniceras* Casey, 1954.

484 Finally, a large number of species are based on material that was collected from condensed
485 beds. This is the case for many key taxa from South-eastern France (Seunes, 1887; Jacob, 1905) and
486 Switzerland (Jacob and Tobler, 1906); and the great majority of the late Aptian faunas from western
487 Caucasus (Egoian, 1965, 1969), northern Caucasus (Sinzow, 1906, 1913; Nikchitch, 1915), Dagestan
488 (Anthula, 1900; Kazansky, 1914), Georgia (Kvantaliani, 1971, 1972), Turkmenistan (Glazunova, 1953;
489 Tovbina, 1968, 1970, 1982) and Mangyschlak (Sinzow, 1908; Glazunova, 1953). Although the recent
490 contribution of Bogdanova and Mikhailova (2016) clarifies the biostratigraphic distribution of the
491 early late Aptian ammonites of the northern Caucasus and Transcaspia to some extent, the precise
492 range of the latest Aptian faunas remain poorly understood.

493 The palaeontological study of the Aptian ammonite faunas from the **EAB** deserve extensive
494 taxonomic descriptions that are beyond the scope of the present paper. Selected elements of the
495 fauna that are crucial for the definition of the biostratigraphic scheme are illustrated (Figs. 8 to 11)
496 and their taxonomic assignments are briefly discussed. In most cases, the identifications are based
497 on direct comparison with the originals or plaster casts of the type material from SE France (Seunes,
498 1887; Jacob, 1905; Kilian and Reboul, 1915), Switzerland (Jacob and Tobler, 1906), Dagestan
499 (Anthula, 1900) and Mexico (Burckhardt, 1925; Humphrey, 1949). A deliberate choice has been
500 made to reduce synonymies to a limited number of key specimens. Emphasis has been taken on
501 material from Morocco previously illustrated in the literature. Unless otherwise mentioned the
502 suprageneric classification retained herein follows the nomenclature of Wright et al. (1996).

503

504 Order Ammonoidea Zittel, 1884

505 Suborder Ancyloceratina Wiedmann, 1966

506 Superfamily Douvilleiceratoidea Parona and Bonarelli, 1897

507 Family Douvilleiceratidae Parona and Bonarelli, 1897

508

509 Comment. For the reasons exposed by Bulot in Vincent et al. (2010, p. 184), Cheloniceratinae Spath,
510 1923 is herein considered as a junior subjective synonym of the subfamily Douvilleiceratinae.

511

512 Subfamily Douvilleiceratinae Parona and Bonarelli, 1897

513 Genus *Procheloniceras* Spath, 1923

514

515 *Type species. Ammonites stobieckii* d'Orbigny, 1850, p. 113.

516

517 *Procheloniceras dechauxi* (Kilian and Reboul, 1915)

518 Figure 8.7–12

519

520 *Holotype. Douvilleiceras Martinii* var. *Dechauxi* Kilian and Reboul, 1915, p. 56, pl. 1, fig. 7, 7b, pl. 7,
521 fig. 2. The specimen is housed in the Université de Grenoble collections (catalogue number UJF-
522 ID.1084).

523

524 *Remarks.* The genus *Procheloniceras* was reported on various occasions from the Tamzergout Fm. of
525 the **EAB**. According to Roch (1930), *Procheloniceras pachystephanum* (Uhlig, 1883) is the most
526 common species with *P. albrechtiaustriae* (Uhlig, 1883). Additionally, *P. stobieckii* (d'Orbigny, 1850)
527 was reported by Rey et al. (1986a, 1988). None of the specimens quoted by those authors were
528 illustrated. The well-preserved and abundant material collected at Tamanar and Zem Zem questions
529 those identifications. Even though a full revision of the genus is needed, the recent contributions of
530 Delanoy (1995, 1998) and Delanoy et al. (2008) has helped to clarify the systematics of
531 *Procheloniceras*. Our material differs from *P. pachystephanum* and *P. albrechtiaustriae* by its smaller
532 umbilicus, more depressed whorl section, and rigid and regular ornamentation. Most Moroccan
533 specimens (Fig 8.7–10) match the holotype of *Procheloniceras dechauxi* well. The variability of the
534 populations includes a slender morphology (Fig. 8.11–12) that superficially matches the lectotype of
535 *P. stobieckii* illustrated by Conte (1981) and Ropolo et al. (2008a).

536

537 Genus *Epicheloniceras* Casey, 1954

538

539 *Type species. Douvilleiceras Tschernyschewi* Sinzow, 1906, by original designation.

540

541 *Epicheloniceras* gr. *subbuxtorfi* – *paucinodum* (Burckhardt, 1925)

542 Figure 9.1–8.

543

544 v 1930. *Douvilleiceras buxtorfi* Jacob and Tobler; Roch, p. 381, pl. 19, fig. 1a–d.

545 v 1930. *Douvilleiceras aequicostatum* Burckhardt; Roch, p. 381, pl. 20, fig. 1a–b.

546 1998. *Chelonicerias (Epicheloniceras) gr. martinoides* Casey; Witam, p. 356, pl. 7, fig. 2–6.

547

548 *Remarks.* *Epicheloniceras* is a fairly common element of the late Aptian ammonite faunas of the **EAB**.

549 It is noteworthy that the large population from Tiskatine (bed **TK 161A**) can easily be distinguished

550 from the *Epicheloniceras* of the *martini* (d'Orbigny, 1841) and *buxtorfi* (Jacob and Tobler, 1906)

551 groups by the early loss of the ventral tubercles, simplified ornamentation and highly distinctive

552 subrounded whorl section. The various morphotypes show close similarities with *Epicheloniceras*

553 *paucinodum* (Burckhardt, 1925), *E. subbuxtorfi* (Burckhardt, 1925) and *E. aequicostatum*

554 (Burckhardt, 1925). This affinity with the Mexican taxa was already recognised by Roch (1930, p.

555 381).

556

557 "*Epicheloniceras*" *marocanus* (Roch, 1930)

558 Figure 9.25–28.

559

560 v 1930. *Parahoplites* (?) *marocanus* Roch, p. 378, pl. 16, fig. 5–6.

561

562 *Lectotype.* *Parahoplites* (?) *marocanus* Roch, 1930, pl. 16, fig. 5 (UJF-ID.1601), herein designated.

563

564 *Type locality.* Tiskatine (= Djebel Tissakatine in Roch, 1930).

565

566 *Description.* Small-sized planulate ammonites ($D_{\max} \leq 50\text{mm}$) with a moderately evolute coiling.

567 Earliest ontogeny unknown. On the adult whorl, the whorl section is depressed, sub-rounded and

568 becomes progressively compressed, sub-rectangular and is higher than it is wide. The venter is

569 moderately flattened and becomes rounded near the aperture. The umbilical wall is rounded and

570 tends to become steep in the adult. The suture line is unknown. Two ornamental stages on the adult
571 whorl:

572 (i) alternation of tuberculate primary and one to two atuberculate secondary ribs. Small
573 rounded tubercles occur on the upper part of the flank and divide into two branches. All ribs cross
574 the venter but the adoral branches bear small thickenings on the ventrolateral margin;

575 (ii) abrupt change toward spaced, simple ribs with rare secondaries. Ribs are slightly flexuous
576 or straight with a marked apertural bending on the upper part of the flank. All ribs cross the venter
577 forming an elevated proverse bending.

578

579 *Remarks.* This micromorphic species is provisionally assigned to *Epicheloniceras*. Derivation from an
580 *Epicheloniceras* stock is based on the similarity of its early ontogenetic stages with the juveniles of
581 the *Epicheloniceras* of the *waageni* (Anthula, 1900) – *tchernyschewi* (Sinzow, 1906) group that
582 occur in underlying level (see Fig. 9.29-32).

583

584 Family Deshayesitidoidea Stoyanow, 1949

585 Family Deshayesitidae Stoyanow, 1949

586 Genus *Deshayesites* Kazansky, 1914

587

588 *Type species.* *Ammonites Deshayesi* d'Orbigny, 1841, by original designation.

589

590 *Deshayesites* aff. *euglyphus* Casey, 1964 in Delanoy (1995)

591 Figure 8.1–6.

592

593 Microconchs

594

595 v 1995. *Deshayesites* aff. *euglyphus* Casey, Delanoy, p. 77, pl. 1, fig. 4, pl. 4, fig. 4.

596 v 1998. *Deshayesites* aff. *euglyphus* Casey, Delanoy, pl. 6, fig. 2, pl. 24, fig. 2 (= Delanoy, 1995, pl. 1
597 ,fig. 4, pl. 4, fig. 4).

598 1998. *Deshayesites* aff. *luppovi* Bogdanova, Witam, p. 357, pl. 8, fig. 2.

599

600 Macroconchs

601

602 v 1995. *Deshayesites* aff. *evolvens* Luppov, Delanoy, p. 77, pl. 3, fig. 1.

603

604 *Remarks.* The Moroccan material includes both microconchs and macroconchs. The microconch
605 forms show an intermediate morphology between *Deshayesites luppovi* Bogdanova, 1983 and the
606 coarser morphotypes of the *D. forbesi* Casey, 1961 group such as *D. euglyphus* Casey, 1964. They
607 match well the specimens illustrated by Delanoy (1995) as *D. aff. euglyphus*. The larger macroconchs
608 compares with specimens from South East France that were most often misidentified as
609 *Deshayesites consobrinus* (d'Orbigny, 1841) (Ropolo et al., 2000a, p. 162-163, fig. 3.2 and 4; Ropolo
610 et al., 2006, pl. 6, fig. 6). The Moroccan specimens are left in open nomenclature and their species
611 assignment will be addressed in a forthcoming contribution once the ongoing revision of the
612 material from South-eastern France is completed.

613

614 Superfamily Acanthohoplitoidea Stoyanow, 1949

615

616 *Remark.* Our ongoing revision of the Acanthohoplitidae supports the view that it has no phyletic link
617 with the Parahoplitidae, even so both families derivate iteratively from the Douvilleiceratidae.

618

619 Family Acanthohoplitidae Stoyanow, 1949

620 Genus *Colombiceras* Spath, 1923

621

622 *Type species. Ammonites crassicostratus* d'Orbigny, 1841, by original designation.

623

624 *Colombiceras tobleri* (Jacob and Tobler, 1906)

625 Fig. 9.9-12

626

627 v 1930. *Parahoplites teffryanus* Karsten in Anthula; Roch, p. 377, pl. 16, fig. 7.

628 ? 1998. *Colombiceras* aff. *tobleri* Jacob and Tobler; Witam, p. 362, pl. 10, fig. 5.

629

630 *Remarks.* Among the material from Morocco two morphologies can be distinguished. The most
631 common is a compressed and finely ribbed morphotype with rectangular section and tabulate
632 venter (Fig. 9.9–10). It somewhat bears a similarity to *Colombiceras crassicostratum* (d'Orbigny, 1841)
633 and *Gargasicerias gargasense* (d'Orbigny, 1841) but the early ontogeny differs by having a distinct
634 ribbing pattern where all the ribs appears single or by pairs on the umbilical margin and cross the
635 venter without weakening. The other morphotype is a coarsely ribbed form that shows a rounded
636 rectangular whorl section higher than wide, and a widely spaced alternation of primary and
637 intercalatory ribs. It closely matches the lectotype of *Colombiceras discooidale* (Sinzow, 1908, pl. 5,
638 fig. 17–18) at similar growth stages. In our opinion the slender and coarse morphotypes are
639 conspecific and are linked by intermediate forms such as the specimen illustrated on Figure 9.9–11.
640 Since the examination of a large number of specimens from South-eastern France has convinced us
641 that *Colombiceras discooidale* represent a compressed morphotype of *C. tobleri*, the Moroccan
642 specimens are considered to fall within the range of variation of the later species. It should be noted
643 that the lectotype of *C. tobleri* notably differs from our material by its rounded section throughout
644 ontogeny.

645

646 Genus *Acanthohoplites* Sinzow, 1908

647

648 *Type species. Parahoplites aschiltaensis* Anthula, 1900, by subsequent designation (Roman, 1938).

649

650 *Remarks.* Re-examination of Anthula's type material leaves no doubt that the lectotype designated
651 by Dimitrova (1967, p. 185) is the microconch of the large specimen illustrated by Anthula (1900, pl.
652 10, fig. 4, pl. 11, fig. 1). Our material includes a series of individuals that match the macroconch well.
653 Even though the large specimen illustrated on Figure 11.1-2 does not show the characteristic
654 juvenile ornamental stage of *A. aschiltaensis*, complete material from other sections support its
655 specific identification. The very large adult body chambers, which are characterized by a rigid adult
656 ornamentation, and were collected from bed **MR 10t** and **11** at Mramer, are left in open
657 nomenclature.

658

659 Genus *Protacanthoplites* Tovbina, 1970

660

661 *Type species. Parahoplites abichi* Anthula, 1900, by original designation.

662

663 *Remarks.* The genus has been variously interpreted since its introduction by Tovbina (1970) and
664 some authors suggest that it could be a junior subjective synonym of *Acanthohoplites* (Wright et al.,
665 1996, Bogdanova and Mikhailova, 2016). Comparison of the lectotypes of *A. aschiltaensis* and *P.*
666 *abichi* shows that the two species are fairly close but the material at our disposal does not allow yet
667 a definitive conclusion regarding synonymy. For the time being, we maintain the genus
668 *Protacanthoplites* to accommodate the Moroccan specimens, that occur above the last occurrence
669 *Acanthohoplites aschiltaensis*, and closely match *P. abichi* (Fig. 9.17–20), *P. bergeroni* (Seunes, 1987)
670 and "*Protacanthoplites*" aff. *multinodosus* Tovbina, 1982 (Fig. 9.13–16).

671

672 Genus *Diadochoceras* Hyatt, 1900

673

674 *Type species. Ammonites nodosocostatus* d'Orbigny, 1841, by original designation.

675

676 *Remarks.* Specimens of *Diadochoceras* from the type locality of *D. nodosocostatum* are fairly
677 numerous in the French historical collections (Obata, 1975), but our understanding of the genus is
678 largely handicapped by the condensed character of faunas and the absence of a proper revision of
679 the type species that is based on a single poorly preserved juvenile specimen (neotype designed by
680 Guerin-Franiette in Gauthier et al., 2006). Examination of the topotype material strongly suggest
681 that the large number of new taxa introduced by Glazunova (1953), Mikhailova (1963), Egoian (1965,
682 1969) and Kvantaliani (1971, 1972) merely represent sexual dimorphism and intraspecific variation.
683 This high variability is reflected by the Moroccan specimens and assignment to a *Diadochoceras*
684 *nodosocostatum* group was preferred to identification of specific typologic taxa (Fig. 9.21-22).

685

686 Genus *Nodosohoplites* Egoian, 1965

687

688 *Type species. Nodosohoplites subplanatus* Egoian, 1965, by original designation.

689

690 *Remarks.* As pointed out by Klein and Bogdanova (2013), there is no agreement in the literature
691 about the validity and content of *Nodosohoplites*. Wright et al. (1996) consider that the genus is a
692 junior subjective synonym of *Diadochoceras*. Szives et al. (2007) suggest that some of the specimens
693 that were placed in *Nodosohoplites* by Egoian (1965) are sexual dimorphs of *Diadochoceras*. Our
694 material shows that the planulate forms with a reduced trituberculate ornamental stage similar to
695 *Nodosohoplites subplanatus* and *N. multispinatus* (Anthula, 1900) appear at a slightly younger
696 stratigraphical level than the first occurrence of *Diadochoceras* of the *nodosocostatum* group.
697 Pending a global revision of *Diadochoceras* and *Nodosohoplites* based on abundant and
698 stratigraphically well-documented populations we prefer to keep the two genera separate.

699

700 Genus *Elsaella* gen. nov.

701

702 *Derivation of name.* Dedicated to Elsa Schnebelen-Bulot for her participation to our field
703 investigations in Morocco and continuous support to one of us (L.G.B.) during the preparation of this
704 work.

705

706 *Type species.* *Elsaella tiskatinensis* sp. nov., by monotypy.

707

708 *Diagnosis.* Small-sized ($D \leq 45\text{--}50$ mm), planulate ammonite with a moderately evolute coiling. Sub-
709 rounded whorl section in the juvenile becomes sub-rectangular as growth increases. Venter is
710 rounded throughout ontogeny. Umbilical wall is low to moderately steep at the adult stage. Two
711 ornamental stage (i) irregular alternation of simple or bifurcate primary ribs and simple secondaries
712 ($D \leq 15$ mm); (ii) Uniform and dense ribs originates by pairs or bundles from periumbilical bullae.
713 Bullae strengthen as growth increases and the ribs are more spaced on the outer part of the body
714 chamber. All ribs cross the venter without interruption and slightly bend forward. Aperture is simple.
715 Acanthohoplite suture line.

716

717 *Discussion.* The genus *Immunitoceras* (type species: *I. immunitum* Stoyanow, 1949) compares
718 superficially to *Elsaella*. The original description and the re-examination of the holotype leave no
719 doubt that the primary ribs are bituberculated at the younger stages and do not compare with the
720 equivalent ontogenetic stages of *Elsaella*. The only other acanthohoplite genus that can be compared
721 with *Elsaella* gen. nov. is *Nolaniceras*. Differences between the type species of the two genera are
722 discussed below.

723

724 *Elsaella tiskatinensis* sp. nov.

725 Fig. 10.1-16.

726

727 1998. *Nolaniceras nolani* Seunes, Witam, p. 361, pl. 9, fig. 7a-b, pl. 10, fig. 1-2 (sol).

728 1998. *Nolaniceras nolani* var. *planulata* Egoian, Witam, p. 361, pl. 10, fig. 3a-b (sol).

729

730 *Holotype*. (MANCH) LL.16123, (Fig. 10.1-2).

731 *Paratypes*. (MANCH) LL.16124 – LL.16131, (Fig. 10.3-16).

732 *Type locality*. Tiskatine, Essaouira-Agadir Basin, Morocco.

733

734 *Type strata*. Bed **TK206**, *Elsaella tiskatinensis* Zone (see definition below), upper Aptian.

735

736 *Diagnosis*. As for the genus.

737

738 *Discussion*. *Elsaella tiskatinensis* sp. nov. has been misidentified as *Nolaniceras nolani* in the

739 Moroccan literature (Rey et al. 1986a, 1988, Witam 1998, Peybernes et al. 2013). In a recent

740 revision, Bulot et al. (2014) have shown that *Nolaniceras nolani* differs from all other uppermost

741 Aptian Acanthohoplitidae by its highly distinctive low and convex umbilical wall and most unusual

742 ornamental style. Compared to *Elsaella tiskatinensis* sp. nov., *Nolaniceras nolani* can be

743 distinguished by its compressed suboval whorl section throughout ontogeny, the absence of

744 umbilical bullae and attenuation of the ribbing on the venter.

745

746 *Remarks*. The late upper Aptian Moroccan faunas also includes a coarser and larger forms of *Elsaella*

747 that compare with *Acanthoceras bigoti* Seunes (1887) and *Acanthoceras migneni* Seunes (1887).

748 These two species have been diversely interpreted in the literature and remain poorly understood

749 since it seems that the originals from the Mignen collection are lost. The coarser forms of *Elsaella*

750 that dominates the assemblages at Mramer and Zem Zem are provisionally referred as *Elsaella* sp. 1

751 and sp. 2. More work is needed to determine if these forms represent ecological morphotypes of *E.*
752 *tiskatinensis* or new taxa.

753

754 *Geographical and stratigraphical distribution.* Since *E. tiskatinensis* is a very common ammonite in
755 the lower part of the late upper Aptian of the **EAB**, it is selected herein as the index species of the *E.*
756 *tiskatinensis* Zone. Outside Morocco, and despite an extensive survey of the literature, none of the
757 specimens previously referred to *Nolaniceras nolani* in the literature can be assigned with certainty
758 to our new species.

759

760 Genus *Hypacanthoplites* Spath, 1923

761

762 *Type species.* *Acanthoceras Milletianum* Var. *plesiotypica* Fritel, 1906, by original designation.

763

764 *Remarks.* Problems with the systematics of the genus *Hypacanthoplites* have been addressed at
765 length by Bulot (2010). The genus was reported on many occasions in the Moroccan literature (Rey
766 et al. 1986a, 1988, Witam 1998, Peybernes et al. 2013). Our new collection shows that
767 hypacanthoplid-like ammonites occur at two different levels in the **EAB**.

768 The older fauna is preserved as small pyritic internal moulds that occur above the *Elsaella*
769 beds at Tiskatine, Tamarar, and Zem Zem. Those forms are provisionally placed in
770 "*Hypacanthoplites*" and left in open nomenclature as sp. 1, sp. 2 and sp. 3. They all show a very clear
771 tabulate venter associated with a weakening of the ribs on the siphonal line. Even though there is a
772 reinforcement of the ribs on both sides on the ventral shoulder; this feature is not as marked as in
773 true *Hypacanthoplites*. The ornamental style of the very finely ribbed "*Hypacanthoplites*" sp. 1 (Fig.
774 11.15-25) suggests evolution from *Elsaella tiskatinensis*. To the difference "*Hypacanthoplites*" sp. 3
775 (Fig. 11.3-11) shows a very distinct bituberculate ontogenetic stage that somehow recalls the

776 ornamentation of *Hypacanthoplites tuberculatus* Egoian, 1969 and *H. microtuberculatus* Egoian,
777 1969.

778 Higher up in the succession, a collection of specimens that match the type material of
779 *Mellegueiceras chihaouiae* Latil, 2011 and "*Hypacanthoplites*" *paucicostatus* Breistroffer in
780 Dubourdieu, 1953 was made. This fauna will be described in a separate paper and its early Albian
781 age is established by the co-occurrence of the genus *Douvilleiceras*. It should be noted that the
782 fragments identified by Witam (1998, pl. 9, fig. 4-6) as *Hypacanthoplites* gr. *jacobi* (Collet, 1907) are
783 merely misidentified "*Hypacanthoplites*" *paucicostatus*.

784

785 **5. Ammonite biostratigraphy and correlation to the Standard Mediterranean Ammonite Scale** 786 **(SMAS)**

787

788 As shown by the systematic notes, the ammonite faunas from the **EAB** have a distinct
789 character that is most certainly linked to the palaeogeography of the basin. Although the successive
790 assemblages are clearly of Tethyan affinity, none of the index species of the Standard Mediterranean
791 Ammonite Scale (**SMAS**) of Reboulet et al. (2011, 2014) were found in the course of our study. As a
792 consequence, we have chosen to introduce a regional biostratigraphic scale based on the main bio-
793 events recognized at the basin scale (Fig. 12). When possible correlations with the **SMAS** are
794 proposed, the precision of these correlations is handicapped by the fact, that for the upper part of
795 Aptian Stage, the biostratigraphic subdivision of the **SMAS** is largely based on assemblage zones and
796 subzones originally defined in the former Soviet Union (Bogdanova and Tovbina, 1995) for which
797 detailed successions are still poorly documented.

798

799 Lower Aptian – *Procheloniceras dechauxi* Zone, new this paper

800

801 *Index species. Procheloniceras dechauxi* (Kilian and Reboul, 1915)

802 *Reference section.* Tiskatine – bed **TK 159** to **TK 160b**.

803

804 *Definition.* The base of the zone is defined by the first occurrence of the index–species that largely
805 dominates the assemblage in all studied sections. The diversity is low but some beds are rich in
806 *Deshayesites* aff. *euglyphus* Casey in Delanoy (1995). It should be noted that *P. dechauxi* and *D.* aff.
807 *euglyphus* are most often mutually exclusive. *Audouliceras* sp. ind., *Pseudohaploceras* cf. *matheroni*
808 (d'Orbigny, 1841) and *Melchiorites melchioris* (Tietze, 1872) are minor elements of the fauna.

809

810 *Correlations.* Correlation with the **SMAS** is based on the occurrence of *P. dechauxi* and *D.* aff.
811 *euglyphus*. These two taxa are known to occur in the lower part of the *Deshayesites forbesi* Zone in
812 South East France (Delanoy, 1995, 1998; Pictet, 2012).

813

814 Upper Aptian – *Colombiceras tobleri* Zone, Eristavi (1960) emended.

815

816 *Index species.* *Colombiceras tobleri* (Jacob and Tobler, 1906)

817 *Reference section.* Tiskatine – bed **TK 161a** to **TK 183c**.

818

819 *Definition.* Since its first introduction, the *Colombiceras tobleri* Zone was variously interpreted in the
820 literature (see discussion in Bogdanova and Mikhailova, 2016). As herein understood, the zone is
821 defined by the first occurrence of its index–species. Two subzones can be distinguished in the **EAB**.

822

823 *Colombiceras tobleri* Subzone, new, this paper

824

825 *Index species.* As for the zone.

826 *Reference section.* Tiskatine – bed **TK 161a** to **TK 163**.

827

828 *Definition.* The base of the subzone is defined by the first occurrence of the index–species. The
829 diversity of the fauna is fairly high. Together with the index species, the various *Epicheloniceras* of
830 the *subbuxtorfi* – *paucinodum* group are the most common elements of the faunal assemblage.
831 Desmoceratids are unusually common compared to the rest of the succession. *Pseudohaploceras*
832 *angladei* Dutour, 2005 non (Sayn, 1891), *Zuercherella zuercheri* (Jacob and Tobler, 1906),
833 *Melchiorites* aff. *emerici* (Raspail, 1831) and *Caseyella* sp. ind. were identified. The heteromorph taxa
834 "*Ammonitoceras*" *lahuseni* (Sinzow, 1906) and ? *Pseudoaustralicerias* sp. are minor elements of the
835 fauna. A single phylloceratid tentatively assigned to *Phylloceras* sp. was collected.

836

837 "*Epicheloniceras*" *marocanus* Subzone, new, this paper

838

839 *Index species.* "*Epicheloniceras*" *marocanus* (Roch, 1930)

840 *Reference section.* Tiskatine – bed **TK 164** to **TK 183c**.

841

842 *Definition.* The base of the subzone is defined by the first occurrence of the index–species. The
843 diversity of the fauna is very low. The lower part of the subzone is marked the acme of
844 "*Epicheloniceras*" *marocanus* and its upper part correspond to a barren interval. The last occurrence
845 of *C. tobleri* is to be noted at the base of the subzone. In the barren interval, spot occurrence of
846 *Epicheloniceras* of the *waageni* (Anthula, 1900) – *tchernyschewi* (Sinzow, 1906) group was
847 identified.

848

849 *Correlations.* Because of the highly endemic character of the fauna, a precise correlation with the
850 **SMAS** is hard to establish. A provisional correlation between the base of the *C. tobleri* Zone of
851 Morocco and the base of the *E. gracile* Subzone (*E. martini* Zone) can be proposed on the basis of
852 the evolution of the genus *Colombiceras* in SE France (Bulot in Dauphin 2002), Caucasus and
853 Transcaspia (Bogdanova and Mikhailova, 2016). Due to the somehow "primitive" character of the

854 Moroccan *C. tobleri* a slightly older position in the *E. martini* Zone cannot be excluded. Ongoing
855 study of the large collection of *Epicheloniceras* should allow more precise correlation in the future.

856

857 Upper Aptian – *Acanthohoplites aschiltaensis* Zone, Mordvilko (1960) emended.

858

859 *Index species.* *Acanthohoplites aschiltaensis* (Anthula, 1900). First

860 *Reference section.* Tiskatine – bed **TK 184** to **TK 205**.

861

862 *Definition.* Since its introduction, the *Acanthohoplites aschiltaensis* Zone has had a number of

863 interpretations in the literature (see discussion in Bogdanova and Mikhailova, 2016). As herein

864 understood the zone is defined by the first occurrence of the index–species. The diversity is very low

865 but a distinctive horizon rich in *Pseudoaustralicerias* of the *ramososeptatum* (Anthula, 1900) group

866 (Fig. 9.23 – 24) was identified in the middle part of the zone. The geographical extension of this bio-

867 event at the scale of the basin remains unknown. In the marginal areas of the **EAB**, the occurrence of

868 large *Epicheloniceras* that do not match any species of the literature is also to be noted. The precise

869 range and affinities of those specimens are still to be documented.

870

871 *Correlations.* The precise range of *A. aschiltaensis* has never been precisely calibrated with the **SMAS**

872 zones. Data from South-eastern France suggests that the lowest occurrence of the *Acanthohoplites*

873 of the *aschiltaensis* group occurs in the uppermost part of the *E. martini* Zone close to its boundary

874 with the *P. melchioris* Zone (Bulot in Dauphin, 2002; Frau and Bulot, unpublished data). According to

875 Russian literature, *A. aschiltaensis* is a common element of the *P. melchioris* Zone (Bogdanova and

876 Mikhailova, 2016, with references).

877

878 Upper Aptian – *Elsaella tiskatinensis* Zone, new this paper

879

880 *Index species.* *Elsaella tiskatinensis* gen. nov. sp. nov.

881 *Reference section.* Tiskatine – bed **TK 206** to **TK 247**.

882

883 *Definition.* The zone is defined by the first occurrence of the genus *Elsaella*. Two subzones can be
884 distinguished.

885

886 *Elsaella tiskatinensis* Subzone, new this paper

887

888 *Index species.* As for the zone.

889 *Reference section.* Tiskatine – bed **TK 206** to **TK 220**.

890

891 *Definition.* The first occurrence of the genus *Elsaella* marks the base of the subzone. In all studied
892 sections the fauna is marked by the evolutive radiation of the Acanthohoplitidae. The various
893 morphologies of *Elsaella* dominate in the lower part of the subzone. In the upper part of the
894 subzone a fairly diverse assemblage of Acanthohoplitidae develops and includes *Diadochoceras* of
895 the *nodosocostatum* group, *Nodosohoplites* of the *subplanatus* group, and various species of
896 *Protacanthoplites*. *Epicheloniceras clansayense* (Jacob, 1905) is a secondary element of the fauna in
897 the marginal parts of the **EAB**.

898

899 "*Hypacanthoplites*" spp. Subzone, new this paper

900

901 *Index species.* Hypacanthoplitids with a tabulate venter ("*Hypacanthoplites*" sp. 1, "*H.*" sp. 2 and "*H.*"
902 sp. 3).

903 *Reference section.* Tiskatine – bed **TK 221** to **TK 247**.

904

905 *Definition.* The fauna is dominated by "*Hypacanthoplites*" sp. 1, "*H.*" sp. 2 and "*H.*" sp. 3 but also
906 contains relict elements from the underlying subzone such as *Protacanthoplites abichi* and
907 *Nodosohoplites gr. subplanatus*. Other noteworthy taxa are *Pseudosilesites seranoniformis* Egoian,
908 1969 (Fig.11.12-14), *Puzosiella minuta* Egoian, 1969 and *Melchiorites ? problematicus* (Fallot and
909 Termier, 1923).

910
911 *Correlations.* The detailed range of the ammonite faunas of the *A. nolani* Zone of the **SMAS** is still
912 not documented. Nevertheless, it seems reasonable to consider that the first occurrence of
913 *Diadochoceras* in the lower part of the *E. tiskatinensis* Zone suggests a correlation with a level close
914 to the base *D. nodosocostatum* Subzone. Co-occurrence of primitive "*Hypacanthoplites*" spp. and
915 *Pseudosilesites seranoniformis* in the upper part of the *A. nolani* Zone was reported from North-
916 eastern Spain by Robert et al. (2001). This suggests that the base of the *Hypacanthoplites* spp.
917 Subzone is likely older than the base of the *H. jacobi* Zone even so correlation with a slightly younger
918 level cannot be excluded.

919
920 In agreement with Latil (2011), the base of the Lower Albian (*M. chihaouiae* Zone) is marked by the
921 co-occurrence of first representatives of the genus *Douvilleiceras* (Fig. 9.33) with *Mellegueiceras*
922 *chihaouiae*. This boundary is clearly documented at Tiskatine and Mramer. At Mramer, the index
923 species of the overlying lower Albian "*Hypacanthoplites*" *paucicostatus* Zone was also identified.

924
925 It should be noted that the beds that contain the first Lower Albian faunas with *Douvilleiceras* (see
926 details below) directly overlie the "*Hypacanthoplites*" spp. Subzone. As a consequence we suspect a
927 hiatus that would embrace the uppermost part of the *H. jacobi* Zone and lower part of the *L.*
928 *tardefurcata* Zone.

929

930 **6. Conclusions**

931

932 High-resolution bed by bed sampling of five sections in the west-central **Essaouira Agadir Basin**933 **(EAB)** has yielded the largest Aptian published ammonite collection in NW Africa. The results provide

934 new information on the age range of previously defined formations, and a type section with fully

935 documented collections correlated against type species. In addition, new species have been

936 identified and the collections provide new insights into the regional endemism of species in this part

937 of NW Africa.

938

939 The main conclusions from the study are:

- 940 • A diverse Aptian and early Albian ammonite fauna is documented, comprising 26 different
941 genera and 43 species.
- 942 • The global palaeobiogeographic character of the fauna is Tethyan, however a high degree of
943 endemism is recognised at the genus and species level.
- 944 • A new genus and species *Elsaella tiskatinensis* is described.
- 945 • New material also allows re-examination of *Parahoplites marocanus*, provisionally included
946 in the genus *Epicheloniceras*.
- 947 • Based on the ammonite distribution a regional biostratigraphic scale is introduced for the
948 Aptian of the **EAB**. 8 zones and subzones are recognised, 5 of which are new.
- 949 • The section at Tiskatine [Lat.: 30.821463° Long.: -9.702555° (Tiskatine 1) and Lat.:
950 30.810477° Long.: -9.739966° (Tiskatine 2)] is selected as the type section for the Aptian in
951 the west-central part of the **EAB**. Key taxa are documented, illustrated with collection
952 references. A correlation based on first occurrence of common taxa with the **Standard**
953 **Mediterranean Ammonite Scale (SMAS)** is proposed when possible.
- 954 • Two basin-scale regional hiatuses are identified. The lower one encompasses the time
955 equivalent of the middle part of the *D. forbesi* Zone to lower part of the *E. martini* Zone of

956 the **SMAS**. The upper one includes the *H. jacobii* Zone and the lowermost part of the *L.*
957 *tardefurcata* Zone equivalent.

958 • The age of the Tamzergout Fm. has been clearly defined, ranging from the local and newly
959 introduced *D. dechauxi* to the *E. tiskatinensis* zones

960 • The top Bouzergoun / base Tamzergout Fm. is identified as being diachronous, ranging in
961 age from the early to late Aptian (*D. dechauxi* to *C. tobleri* zones)

962 • The improved age dating of the Lemgo Fm. (based on collection from the Mramer section)
963 indicates a range from middle upper Aptian *A. ashiltaensis* Zone to the lowermost Albian
964 "*H.*" *paucicostatus* Zone

965 • An early Albian age for the base of the Oued Tidzi Fm. is established by the lowest
966 occurrence of the genus *Douvilleiceras*

967

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1523 Captions ammonite figures

1524

1525 Figure 1. Digital elevation model with sub-crop of geological maps, showing Lower Cretaceous
1526 outcrops of the Essaouira Agadir Basin, studied sections and key locations.

1527

1528 Figure 2. Generalised lithostratigraphy of the upper Bouzergoun to lower Oued Tidzi formations, and
1529 depositional environments in the west-central part of the EAB.

1530

1531 Figure 3. Tiskatine section. Ammonoid distribution and biostratigraphic interpretation.

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1533 Figure 4. Tamanar section. Ammonoid distribution and biostratigraphic interpretation.

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1535 Figure 5. Zem Zem section. Ammonoid distribution and biostratigraphic interpretation.

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1537 Figure 6. Oued Tlit section. Ammonoid distribution and biostratigraphic interpretation.

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1539 Figure 7. Mramer section. Ammonoid distribution and biostratigraphic interpretation.

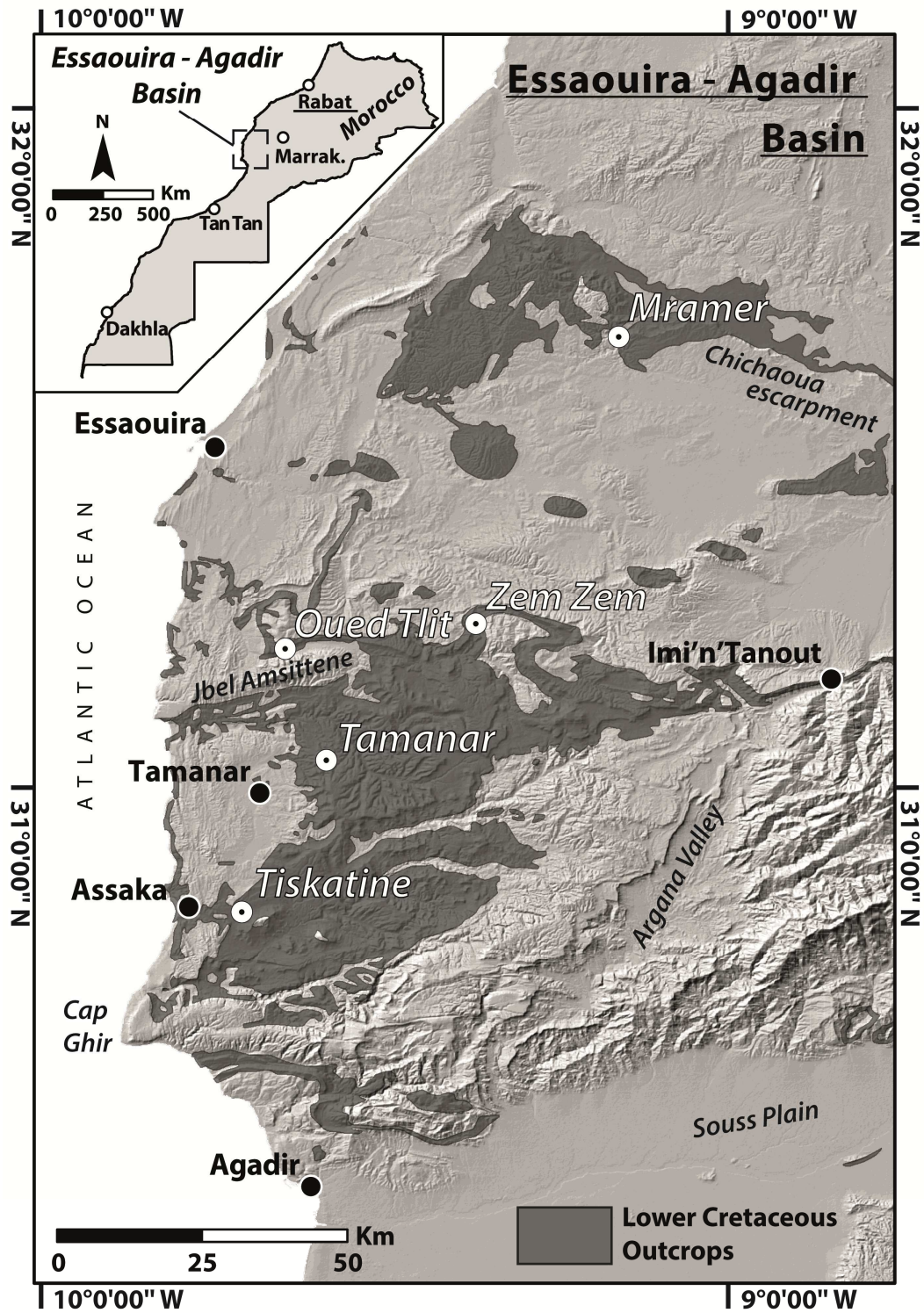
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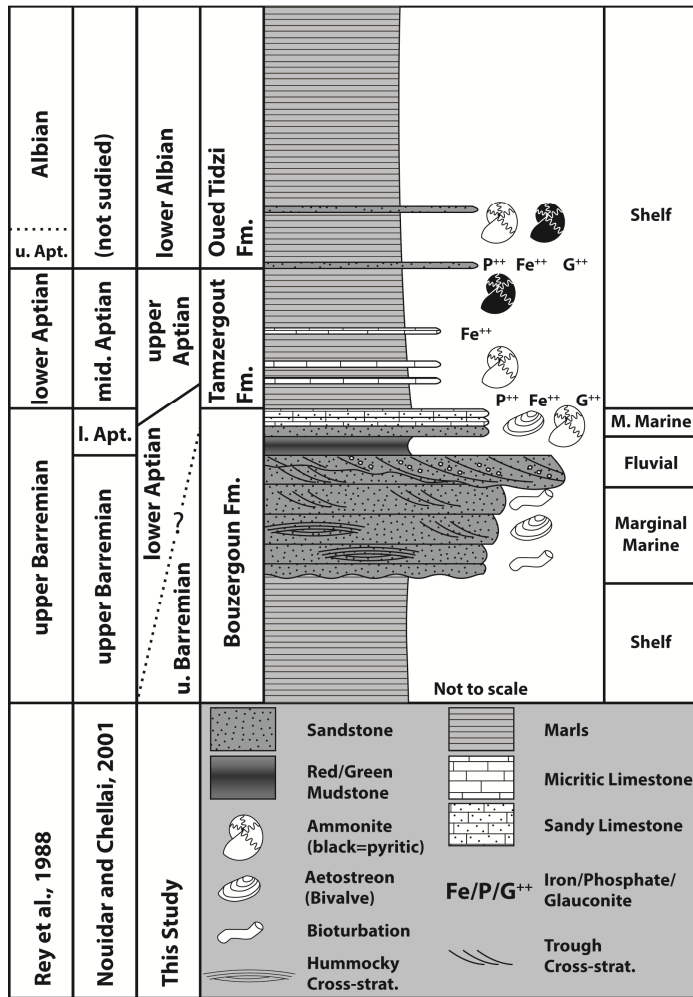
1541 Figure 8. (1–6) *Deshayesites* aff. *euglyphus* (Casey, 1964) in Delanoy (1995) – (1–2) and (5–6) from
1542 bed **TK 160** ((MANCH) LL.16103 and (MANCH) LL.16105), (3–4) from bed **TR 70** (MANCH) LL.16104;
1543 (7–12) *Procheloniceras dechauxi* (Kilian and Reboul, 1915) from bed **TR 69** ((MANCH) LL.16106 –
1544 (MANCH) LL.16108). All specimens coated with ammonium chloride prior to photography. Scale bar
1545 is 10mm.

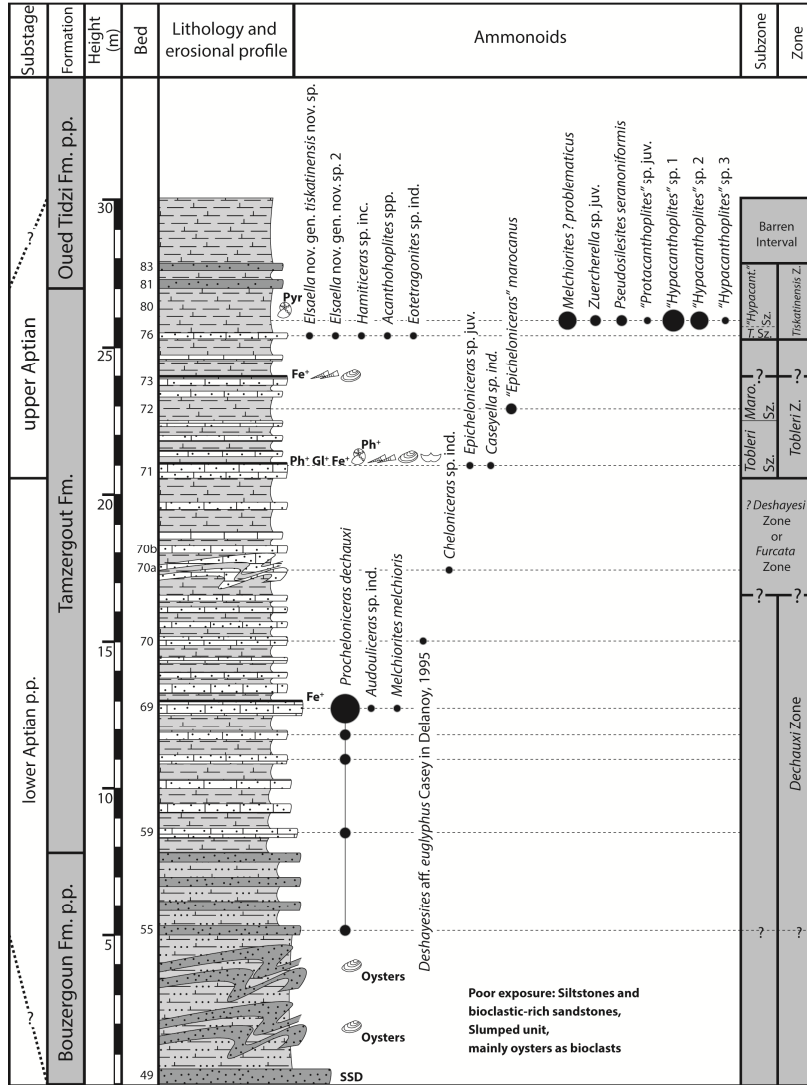
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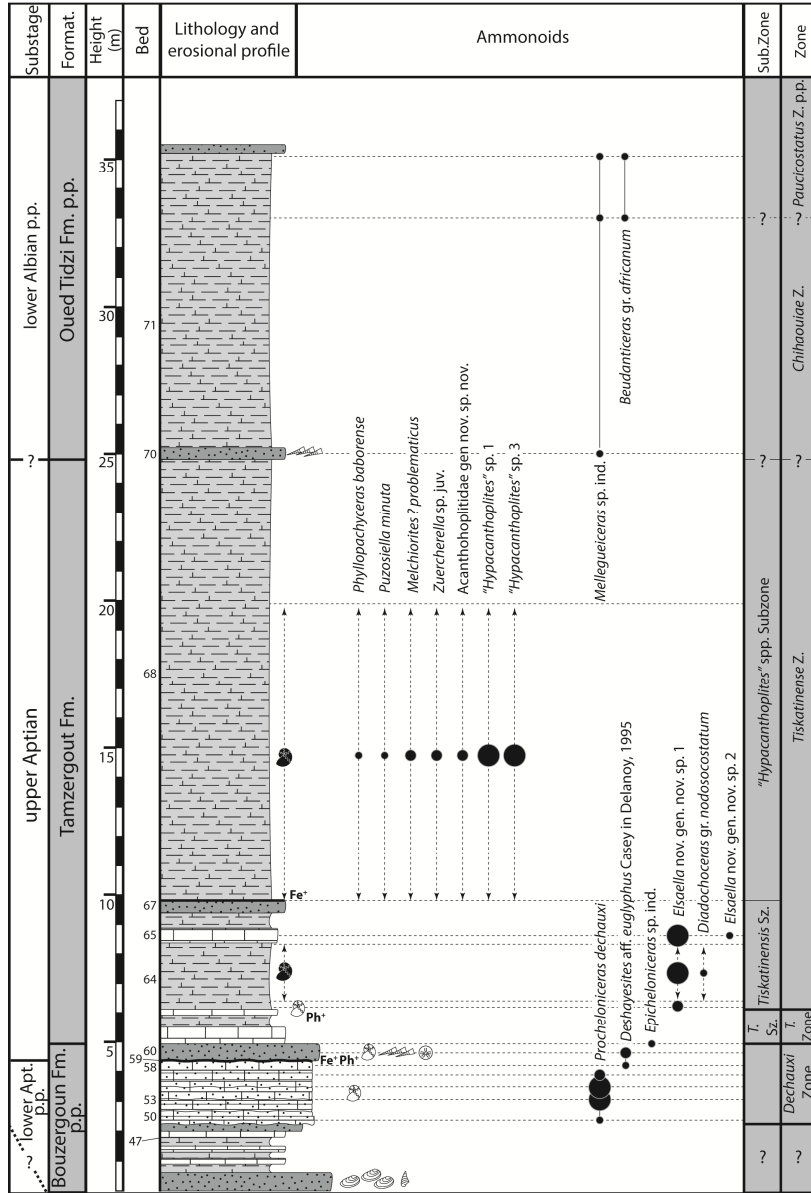
1547 Figure 9. (1–8) *Epicheloniceras* gr. *subbuxtorfi* – *paucinodum* (Burckardt, 1925) from bed **TK 161A**
1548 ((MANCH) LL.16109 – (MANCH) LL.16111); (9–12) *Colombiceras tobleri* (Jacob and Tobler, 1906) from
1549 bed **TK 161A** ((MANCH) LL.16112 and (MANCH) LL.16113); (13–16) "*Protacanthoplites*" aff.
1550 *multinodosus* (Tovbina, 1982) from bed **TK 206** ((MANCH) LL.16114); (17–20) *Protacanthoplites*
1551 *abichi* (Anthula, 1900) from **TK 212/213** ((MANCH) LL.16115); (21–22) *Diadochoceras* gr.
1552 *nodosocostatum* (d'Orbigny, 1841) from bed **TK 206**((MANCH) LL.16116); (23–24)
1553 *Pseudoaustraliceras* gr. *ramososeptatum* (Anthula, 1900) from **TK 196** ((MANCH) LL.16117); (25–28)

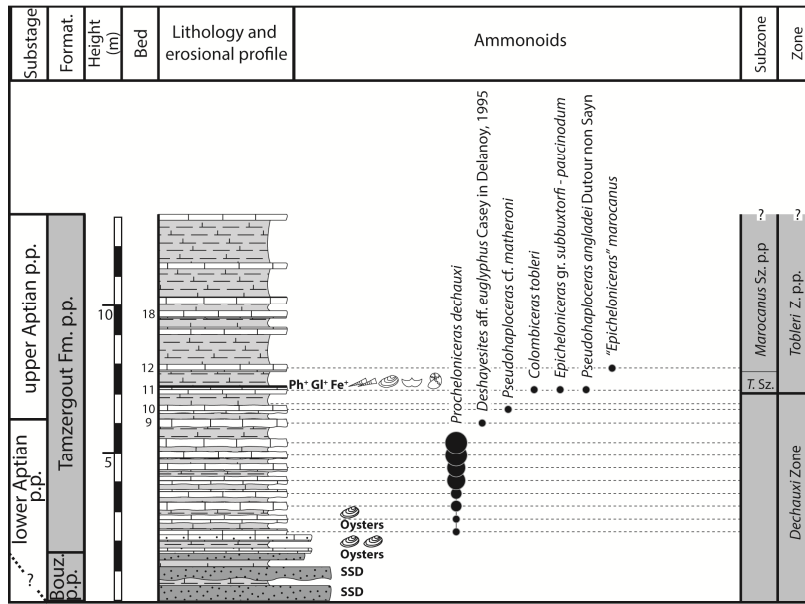
- 1554 "*Epicheloniceras*" *marocanus* (Roch, 1930) from bed **TK 166** ((MANCH) LL.16118 and (MANCH)
1555 LL.16119); (29–32) *Epicheloniceras* sp. juv. 1 gr. *waageni* (Anthula, 1900) – *tschernyschewi* (Sinzow,
1556 1906) from bed **TK 163** (MANCH) LL.16120 and (MANCH) LL.16121); (33) *Douvilleiceras* cf.
1557 *leightonense* (Casey, 1962) from bed **TK 248** ((MANCH) LL.16122). All specimens coated with
1558 ammonium chloride prior to photography. Scale bar is 10mm.
- 1559
- 1560 Figure 10. *Elsaella tiskatinensis* gen. and sp. nov. from bed **TK 206** – (1–2) holotype ((MANCH)
1561 LL.16123), (3–16) paratypes (MANCH) LL.16124 – (MANCH) LL.16131). All specimens coated with
1562 ammonium chloride prior to photography. Scale bar is 10mm.
- 1563
- 1564 Figure 11. (1–2) *Acanthohoplites aschiltaensis* (Anthula, 1900) from bed **TK 192** ((MANCH) LL.16132);
1565 (3–11) "*Hypacanthoplites*" sp. 3 – (3–8) are from bed **ZZ 68** ((MANCH) LL.16133 and (MANCH)
1566 LL.16134), (9–11) from bed **TM 80** ((MANCH) LL.16135); (12–14); *Pseudosilesites seranoniformis*
1567 (Egoian, 1969) from bed **TM 80** ((MANCH) LL.16136); (15–25) "*Hypacanthoplites*" sp. 1 from bed **TM**
1568 **80** ((MANCH) LL.16137 – (MANCH) LL.16140). All specimens coated with ammonium chloride prior to
1569 photography. Scale bar is 10mm.
- 1570
- 1571 Figure 12. Chart showing correlation between the **EAB** and **SMAS** ammonite scales (**SMAS** zones and
1572 subzones after Reboulet et al. 2011, 2014).

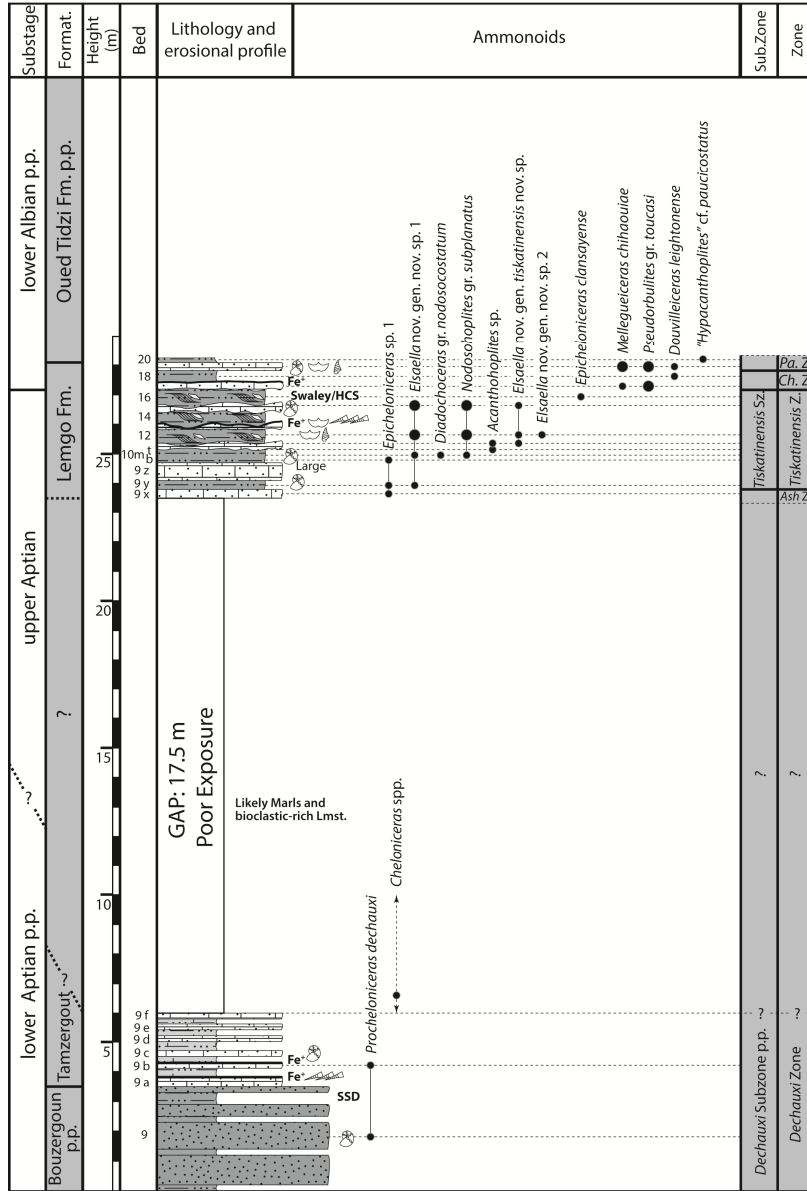


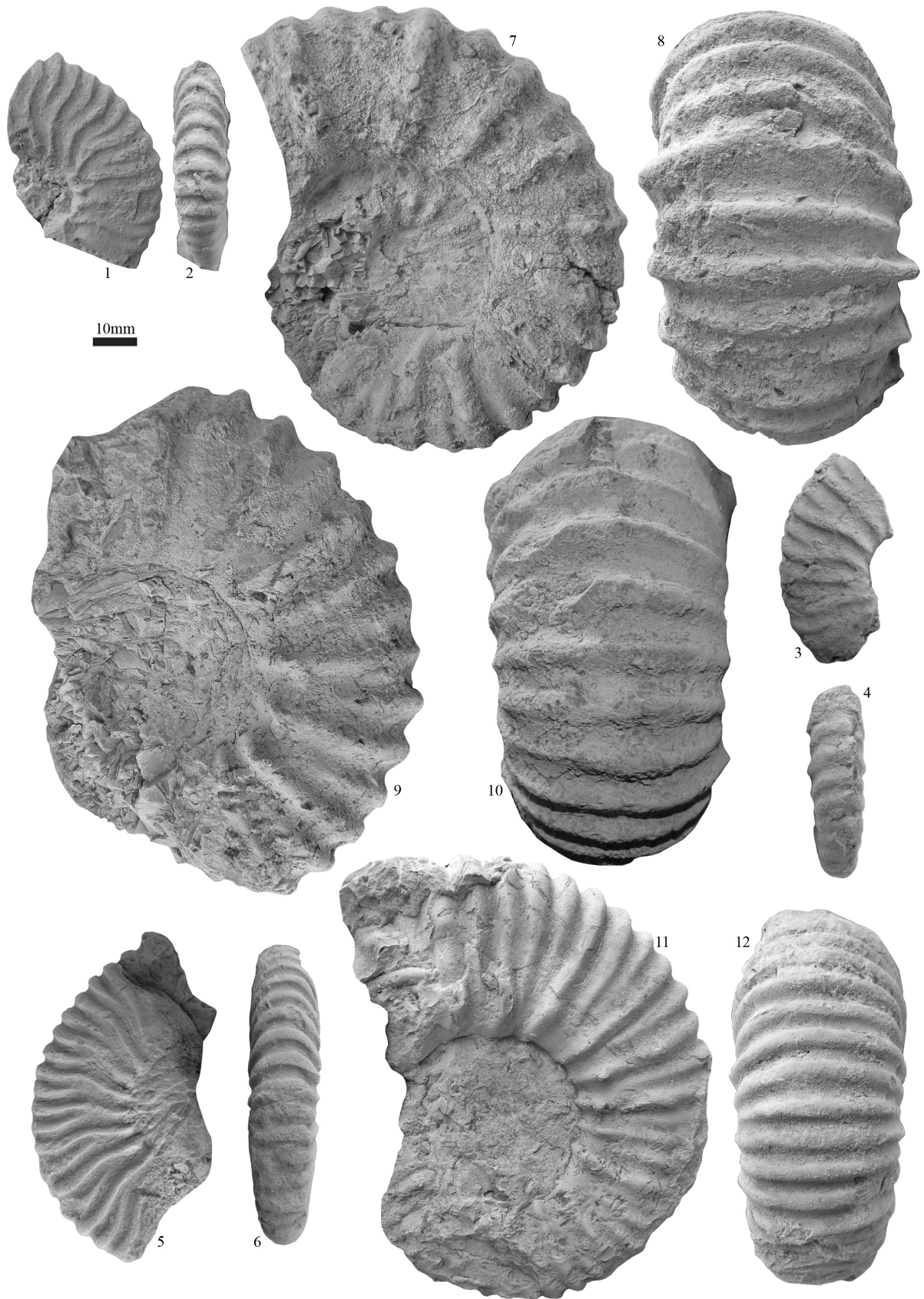


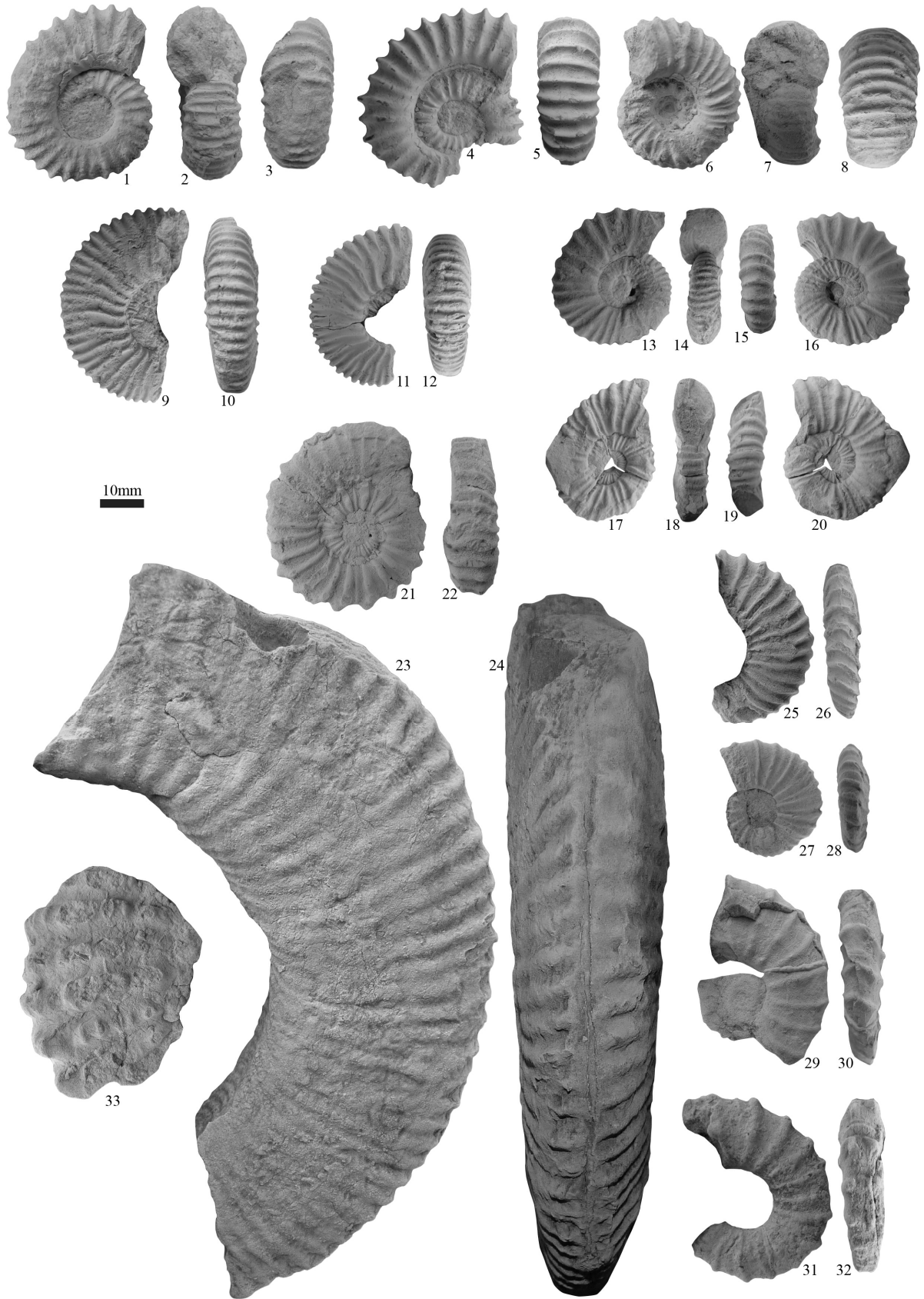


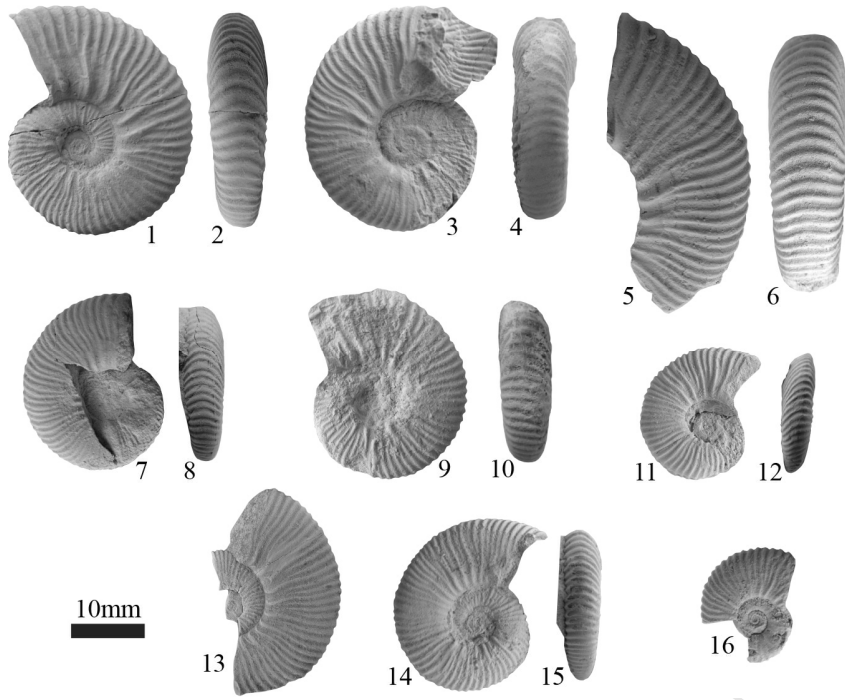


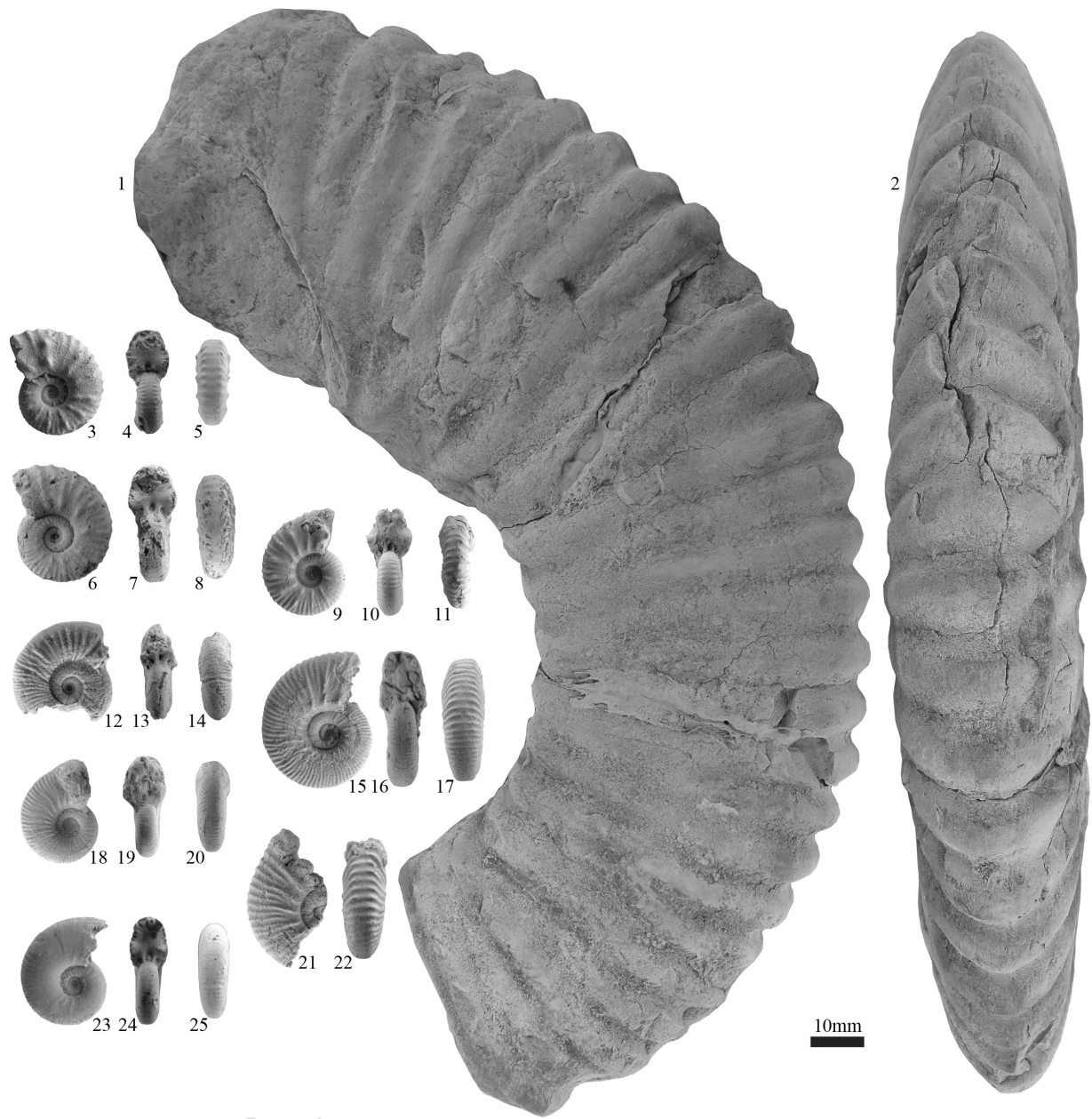












ACC

Stages		Zones This study EAB	Subzones This study EAB	Key Bioevents	Subzones	Zones	Stages	
Albian	lower Alb. p.p.	<i>Mellegueiceras chihauouiae</i>		↑ <i>Douvilleiceras</i> ↑		<i>Leymeriella tardefurcata</i>	lower Alb. p.p.	
		Suspected Hiatus						<i>Hypacanthoplites jacobi</i>
Aptian	upper	<i>Elsaiella tiskatinensis</i>	" <i>Hypacanthoplites</i> " spp. <i>Elsaiella tiskatinensis</i>	↑ " <i>Hypacanthoplites</i> " ↑	<i>Diadochoceras nodosocostatum</i>	<i>Acanthohoplites nolani</i>	upper	
		<i>Acanthohoplites ashiltaensis</i>		↑ <i>Nodosohoplites</i> ↑		<i>Parahoplites melchioris</i>		
		<i>Colombiceras Tobleri</i>	Barren Interval	↑ <i>Acanthohoplites</i> ↑	<i>Epicheloniceras buxtorfi</i>	<i>Epicheloniceras martini</i>		
			" <i>Epicheloniceras</i> " maroccanus		<i>Epicheloniceras gracile</i>			
			<i>Colombiceras tobleri</i>	↑ <i>C. tobleri</i> ↑	<i>Epicheloniceras debile</i>			
	lower	Hiatus				<i>Dufrenoyia dufrenoyi</i>	<i>Dufrenoyia furcata</i>	lower
		Slumps				<i>Dufrenoyia furcata</i>		
						<i>Deshayesites grandis</i>	<i>Deshayesites deshayesi</i>	
						<i>Roloboceras hambrovi</i>		
		<i>Procheloniceras dechauxi</i>		↑ <i>D. aff. euglyphus</i> ↑ ↑ <i>P. dechauxi</i> ↑			<i>Deshayesites forbesi</i>	
	?	?		<i>Deshayesites luppovi</i>	<i>Deshayesites oglanlensis</i>			