

Insights into soft-part preservation from the Early Ordovician Fezouata Biota

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Abstract :

The Fezouata Biota in Morocco is the only Lower Ordovician Lagerstätte yielding a biologically diverse assemblage in a fully marine environment, whilst also containing organisms typical of Cambrian Burgess Shale-type (BST) ecosystems. Fossils from the Fezouata Shale share the same mode of preservation as Cambrian BST biotas defined by carbonaceous compressions and accessory authigenic mineralization. Most organisms of the Fezouata Biota were already dead and decaying on the seafloor when they were buried in-situ by occasional storm-induced deposits in an environment just below the storm-weather wave base. Pre-burial decay in the Fezouata Shale was responsible for the non-preservation of completely cellular organisms such as jellyfish. These conditions contrast with the processes described for soft-tissue preservation in the Burgess Shale (Canada) and the Chengjiang Biota (China). In these two Cambrian Lagerstätten, animals were transported alive or shortly after death by obrution events to an environment that was favorable for preservation. Despite preservational biases, the autochthonous assemblages of the Fezouata Shale offer a unique opportunity to decipher the structure of in situ communities and ecological dynamics in Early Palaeozoic seas, when compared to the allochthonous communities of most Cambrian BST biotas.

Keywords : Exceptional preservation, Cambrian, Fezouata Shale, Burgess Shale, Chengjiang Biota, Taphonomy

44 **1. Introduction**

45 Fossils are key elements in deciphering ancient life on Earth. Much of our knowledge on
46 biodiversification and extinction events comes from mineralized parts such as bones and
47 shells because these are relatively abundant and are commonly found around the globe (Fan et
48 al., 2020). However, organisms having mineralized parts constituting at least part of their
49 bodies are not the sole components in modern ecosystems. A large number of animals are
50 completely soft, having either cuticularized body walls (i.e. formed of polysaccharides), such
51 as annelids and priapulids, or even entirely cellular bodies, such as jellyfish (e.g. Liu et al.,
52 2008; Zhang et al., 2008; Lamsdell et al., 2013; Duan et al., 2014; Lei et al., 2014; Gutiérrez-
53 Marco and García-Bellido, 2015; Martin et al., 2016a; Lerosey-Aubril et al., 2017). Thus,
54 studies based on mineralized parts in the fossil record provide incomplete samples of past
55 animal life on Earth. For this reason, incorporating information from localities with
56 exceptional fossil preservation yielding labile anatomies is crucial to reconstruct ancient
57 ecosystems more accurately (Daley et al., 2018). Although generally rare over the geological
58 time scale, exceptionally preserved biotas discovered in deposits called “Lagerstätten” are
59 common in the Cambrian (Gaines, 2014). The most famous Cambrian site with exceptional
60 preservation is the Burgess Shale (Miaolingian, Canada; Butterfield 1990, 1995; Conway
61 Morris, 1992; Gaines, 2014). The discovery of soft animal taxa in this locality transformed
62 palaeontological knowledge on the earliest eumetazoan-dominated communities during the
63 Cambrian Explosion (e.g. Daley et al., 2009, 2018; Smith and Caron, 2010; Moysiuk et al.,
64 2017; Moysiuk and Caron, 2019; Nanglu et al., 2020). In the last 40 years, over 50 Burgess
65 Shale-type (BST) assemblages have been discovered, most of them from the early to middle
66 Cambrian (Gaines, 2014). Fossils from the Chengjiang Biota (Cambrian Series 2, China)
67 preserved tissues that decay fast in laboratory conditions, and shed light on the early evolution
68 of numerous metazoan phyla (Hou et al., 2004). For instance, nervous tissues were discovered

69 in different arthropod groups ending long-standing debates on the systematic affinities of
70 these taxa (Ma et al., 2012, 2015; Tanaka et al., 2013; Cong et al., 2014). The Chengjiang
71 Biota yielded also the best fossilized cardiovascular system ever discovered (Ma et al., 2014).
72 All these animals from Cambrian BST assemblages were preserved under similar
73 environmental conditions and share the same mode of preservation (Gaines et al., 2008,
74 2012a). They were transported from their living environment, alive or shortly after their death
75 by obrution events, to another setting, where they were buried and eventually preserved
76 (Gaines, 2014). The rapid transport and burial of these animals provided a very limited time
77 for oxic decay to occur, and increased the chances of tissues to escape oxygen in the water
78 column of their original environment (Gaines, 2014). In the deeper facies where they were
79 buried, anoxia was permissive at least at the sea bottom, and carbonate cement precipitated on
80 top of burial event deposits blocking exchange between the water column and sediments and
81 inhibiting oxidants from attaining decaying carcasses (Gaines et al., 2012a). It was also
82 recently suggested that specific clay minerals may have helped BST preservation by slowing
83 down bacterial decay (McMahon et al., 2016). Thus, carcasses were isolated in a fine-grained
84 lithology allowing their preservation in minute details as carbonaceous compressions (Gaines
85 et al., 2008). In some cases, authigenic mineralization (i.e. pyritization, phosphatization) may
86 have occurred, but this remained accessory to the primary carbonaceous mode of preservation
87 (Gaines et al., 2008). Then, for some sites (i.e. the Burgess Shale, and Sirius Passet), the
88 compressed organic matter was kerogenized and matured under metamorphic conditions at
89 temperatures between 300 and 400 degrees (Topper et al., 2018). Although the general
90 conditions for exceptional fossil preservation are relatively well-known for Cambrian
91 Lagerstätten, the mechanisms at play for soft-tissue preservation in younger BST deposits
92 remain largely unexplored.

93 In the early 2000s, a new Lagerstätte was discovered in the Zagora area, Central Anti-Atlas of
94 Morocco. The Fezouata Shale is, so far, the only unit to yield a highly diverse, fully marine
95 exceptionally preserved Ordovician biota (Van Roy et al., 2010, 2015a; Lefebvre et al.,
96 2016a; Martin et al., 2016a). With more than 185 taxa of marine invertebrates recovered from
97 numerous sites in the Zagora area, this formation offers new insights into the diversification
98 of metazoans, at a key interval between the Cambrian Explosion and the Ordovician
99 Radiation (Van Roy et al., 2010, 2015a; Lefebvre et al., 2016a). Most of these taxa are shelly
100 organisms typical of the Ordovician Radiation including asterozoans, bivalves,
101 rhynchonelliformean brachiopods, cephalopods, crinoids, gastropods, graptolites, ostracods,
102 and trilobites (Fig. 1). The Fezouata Biota also comprises a high number of soft-bodied to
103 lightly sclerotized taxa (Fig. 1). Some of these exceptionally preserved organisms (e.g.
104 cirriped crustaceans, eurypterid and xiphosuran chelicerates) represent the oldest occurrences
105 of important marine invertebrate groups, previously only recorded from younger Lagerstätten
106 (Van Roy et al., 2015a). The Fezouata Biota also includes abundant representatives of taxa
107 typical of Cambrian age BST Lagerstätten (e.g. radiodonts, protomonaxonids, armored
108 lobopodians, marrellomorphs, naraoiids) (Botting, 2007, 2016; Vinther et al., 2008, 2017;
109 Van Roy et al., 2010, 2015b; Pérez-Peris et al., in press).

110 Two modes of exceptional preservation have been documented in the Fezouata Shale. The
111 first one occurs in concretions (Gaines et al., 2012b). The second type of exceptional
112 preservation is associated with shales in a generally shallower environment in comparison to
113 the classical Burgess Shale (Martin et al., 2016a; Vaucher et al., 2016). Most Fezouata BST
114 fossils collected in shales are preserved as molds or imprints on the sediments (Martin et al.,
115 2016a), and it is unclear whether these organisms were originally preserved as carbonaceous
116 compressions. In some cases, some non-biomineralized tissues in flattened fossils, such as

117 trilobite digestive tracts and echinoderm water-vascular systems, are preserved in 3D iron
118 oxides (Van Roy et al., 2010; Gutiérrez-Marco et al., 2017; Lefebvre et al., 2019).

119 Considering that numerous mechanisms may favor or alter the preservation of original
120 anatomies in fossils (Fig. 2), deciphering the taphonomic processes is essential for
121 palaeontological and ecological studies, especially for taxa without extant representatives.
122 Consequently, the aim of this study is to review soft tissue taphonomy in the Fezouata Shale
123 based on a multidisciplinary approach combining data across palaeontology, sedimentology,
124 geochemistry and mineralogy. This in-depth reconstruction starts at the life of an organism in
125 its environment and ends at its discovery in surface sediments passing through biostratinomy,
126 diagenesis, metamorphism, and modern weathering (Fig. 2) (Sansom et al., 2010; Bath
127 Enright, 2018; Parry et al., 2018; Purnell et al., 2018). To do so, we will answer the five
128 following questions:

- 129 • Question 1: What is the stratigraphic distribution of exceptional preservation in the
130 Fezouata Shale?
- 131 • Question 2: What were the sedimentary surface processes affecting organisms?
- 132 • Question 3: Under what conditions did decay and diagenetic mineralization take
133 place?
- 134 • Question 4: What were the post-diagenetic processes?
- 135 • Question 5: What is the fidelity of preservation in the Fezouata Shale?

136 Answering these questions allows for comparison of the Fezouata Shale communities and
137 their preservation with both the Burgess Shale and the Chengjiang Biota. This comparison is
138 essential to constrain preservation biases within exceptionally preserved biotas and thus to
139 reconstruct early animal ecosystem evolution. This work has implications in understanding
140 the earliest radiations of complex metazoans on Earth from a fresh perspective that accounts
141 for individual preservation biases that were likely operational at each site.

142

143 **2. Question 1: stratigraphic context – discontinuous occurrences of exceptional**
144 **preservation**

145 The Fezouata Shale is largely exposed in the Anti-Atlas of Morocco. During the Early
146 Ordovician, this area was located at high latitudes close to the palaeo-South pole (Torsvik and
147 Cocks, 2011, 2013). In the Anti-Atlas, the Ordovician succession (maximum ~2500m thick to
148 the West) was originally divided into four lithostratigraphic groups, which are in stratigraphic
149 order: the Outer Feijas, the First Bani, the Ktaoua, and the Second Bani (Fig. 3) (Choubert,
150 1942). Following the stratigraphic work of Destombes in the second half of the 20th century,
151 these stratigraphic groups were subdivided into several formations (Fig. 3) (Destombes, 1970,
152 1971; Destombes et al., 1985). The Fezouata Shale (Tremadocian–Floian) is unconformably
153 deposited over the middle Cambrian Tabanite Group and is conformably overlain by the
154 Zini Formation (late Floian) that is itself overlain by the Tachilla Formation (Darriwilian)
155 (Fig. 3) (Destombes, 1970, 1971; Destombes et al., 1985). The Fezouata Shale is a siltstone-
156 dominated formation outcropping in an area of 900 km² in the Zagora region and with a total
157 thickness of ~ 900 m (Martin et al., 2016b, Vaucher et al., 2016) with sandstone layers
158 becoming more common in the upper part of the formation (Vaucher et al., 2016, 2017).
159 Although mineralized fossils were discovered since the early excavations in the first half of
160 the 20th century, exceptional fossil preservation in the Fezouata Shale was not documented
161 until the early years of the 21st century (Van Roy et al., 2010; Lefebvre et al., 2016a). In the
162 Fezouata Shale, the distribution of exceptional preservation is not random. Exceptionally
163 preserved fossils are found in two distinct stratigraphic intervals (Lefebvre et al., 2016a,
164 2018). Based on acritarchs, conodonts, and graptolites (Gutiérrez-Marco and Martin, 2016;
165 Lehnert et al., 2016; Nowak et al., 2016; Lefebvre et al., 2018), a late Tremadocian age (Tr3)
166 was proposed for the lower, about 70-m thick interval (*A. murrayi* graptolite biozone; Fig. 3).

167 The upper interval is thinner (~50-m thick), and it occurs about 240 m higher in the
168 succession (Lefebvre et al., 2018). Graptolites suggest a mid-Floian age (Fl2) for this upper
169 interval (Fig. 3) (Gutiérrez-Marco and Martin, 2016; Lefebvre et al., 2018). This review
170 focuses on exceptionally preserved material from the lower interval, because it is
171 stratigraphically well constrained (Van Roy et al., 2010; Martin et al., 2016a; Lefebvre et al.,
172 2018). Within this interval, soft-parts are not found everywhere (Saleh et al., 2019). They
173 occur in discontinuous levels with a periodicity of 100,000 years pointing to a possible
174 eccentricity control through seasonality on this type of preservation (Fig. 3) (Saleh et al.,
175 2019). The control of eccentricity on seasonality (Fig. 3) suggests that the conditions favoring
176 this type of preservation were more likely to be ephemeral than permanent (Saleh et al.,
177 2019).

178

179 **3. Question 2: surface processes – *in-situ* burial by distal tempestites**

180 The lithology of the Fezouata Shale varies from claystone to fine-grained sandstone (Vaucher
181 et al., 2016, 2017; Saleh et al., 2020a), while most of the sedimentary succession of this
182 formation is constituted of siltstones (Vaucher et al., 2016, 2017; Saleh et al., 2020a). The
183 sedimentary structures found in the Fezouata Shale are typical of a storm-wave dominated
184 environment (Fig. 4A) (Martin et al., 2016; Vaucher et al., 2016, 2017; Saleh et al., 2020a).
185 These storms were erosive, and their deposits are characterized by the presence of normally
186 graded beds (Fig. 4A, C) (Saleh et al., 2020a), and by oscillatory ripples that were increasing
187 in wavelength from distal (null to millimetric wavelength) to proximal settings (metric to
188 plurimetric wavelength) (Fig. 4B, D) (Vaucher et al., 2016, 2017). Storm-wave deposits are
189 discontinuous in the Fezouata Shale. In proximal settings, a tidal modulation of storm waves
190 occurred and is recorded as a repeated stack of larger (low tide) to smaller (high tide)
191 oscillatory structures within fine-grained sandstones (Fig. 4E) (Vaucher et al., 2017). A

192 statistical correlation between sedimentological data from cores and palaeontological data
193 from outcrops showed that exceptional preservation is associated with one of the most distal
194 facies of the Fezouata Shale (Fig. 5) (Saleh et al., 2020a). This facies is characterized with an
195 abundance of mudstones (i.e. an average of 60% illite
196 $\{(K,H_3O)(Al,Mg,Fe)_2(Si,Al)_4O_{10}[(OH)_2,(H_2O)]\}$, 10% of various chlorite minerals, and 30%
197 siltstones (quartz SiO_2) (Saleh et al., 2019). In this facies, event deposits are not stacked and
198 are separated by background sediments (Saleh et al., 2020a). Furthermore, oscillatory
199 structures are absent indicating that these sediments represent storm-induced deposits in an
200 environment just below the storm wave base (SWB; Fig. 5). Furthermore, exceptionally
201 preserved fossils in this facies are interpreted as autochthonous, because they are occurring
202 right under and not within burial deposits (Fig. 4F, G) (Vaucher et al., 2016; Saleh et al.,
203 2018; 2020a,b). When fossils are disarticulated, this results from in-situ decay rather than
204 transportation, because they do not show any preferential orientation (Fig. 4F) with little
205 evidence of physical abrasion (Martin et al., 2015; Lefebvre et al., 2016a; Saleh et al., 2018;
206 Vannier et al., 2019). In fact, only strong storms are accountable for the accumulation of very
207 coarse siltstones to very fine sandstones in this setting causing an entombment delay and the
208 decay of dead organisms on the seafloor (Saleh et al., 2020a). Due to this pre-burial decay,
209 many fossiliferous intervals yielded hundreds of fossils from which only tens preserved soft-
210 structures (Lefebvre et al., 2019; Saleh et al., 2020a). The rarity of storm-induced deposits in
211 distal settings and its impact on living communities in the Fezouata Shale is validated through
212 observations on body-size fluctuations between sites of this formation (Saleh et al., 2018;
213 2020b). In proximal sites, sessile epibenthic organisms were recurrently buried by storm
214 deposits and could not attain large sizes (Fig. 5). On the contrary nearshore endobenthic taxa
215 were not affected by these events and, if sufficiently motile, they could survive and reach
216 larger sizes (Fig. 5) (Saleh et al., 2018). It was also evidenced that vagile benthic trilobite

217 taxa, distributed in all fossiliferous environments of the Fezouata Shale, reached larger size in
218 distal settings of this formation due to minimal physical stress in distal environments (Fig. 5)
219 (Saleh et al., 2020b). Some of them possibly migrated during storm seasons from proximal to
220 distal settings (Vannier et al., 2019). However, an increase in body-size for epibenthic taxa
221 from proximal to distal sites of the Fezouata Shale should not be generalized. Specific levels
222 in intermediate settings of this formation are characterized by low diversity assemblages with
223 an abundance of small-sized individuals (Fig. 5) (Martin, 2016; Martin et al., 2016b). This
224 possibly indicates that oxygenation was not stable and periods with lower oxygen
225 concentration existed in these settings, suggesting a temporary or seasonal oxygen minimum
226 zone (OMZ)-like conditions (Fig. 5). This hypothesis needs further testing using a
227 geochemical approach.

228

229 **4. Question 3: early diagenesis – controlled decay and authigenic mineralization**

230 In the Fezouata Shale there must be some conditions controlling pre-burial decay and
231 compensating for the delay in fossil entombment for exceptional preservation to occur. Pre-
232 burial decay in the Fezouata Shale was controlled and slowed down by a favorable clay
233 mineralogy (Saleh et al., 2019). Chamosite (i.e. originally berthierine) (Fig. 6C) appears to be
234 present in all analyzed levels recording exceptional preservation (i.e. 6 in total), and absent
235 from intervals with skeletal preservation (i.e. 7 levels) (Saleh et al., 2019). Both berthierine
236 and its primary precursor (Anderson et al., 2018) have been shown to slow down decay under
237 experimental conditions with open oxygenic atmospheric diffusion by damaging bacterial
238 cells (McMahon et al., 2016). This may be the main factor that helped some labile anatomies
239 survive delays in entombment by storm-induced deposits in distal facies (Saleh et al., 2019).
240 When burial occurs, and if it leads to the establishment of anoxic conditions, another type of
241 decay takes place (Fig. 2). Anoxic decay transforms organic matter from decaying carcasses

242 with sulfates SO_4^{2-} from seawater into sulfides H_2S . SO_4^{2-} is not a limiting parameter for this
243 reaction in marine environments. Thus, the H_2S output is mainly controlled by the decay
244 products of biological tissues. High quantities of available organic material such as in the
245 cases of giant decaying arthropods (e.g. 2 meters long *Aegirocassis* radiodonts) leads to the
246 establishment of prominent chemical gradients around carcasses and to the early precipitation
247 and mineral overgrowth around tissues, resulting in their preservation in concretions (Gaines
248 et al., 2012b). However, normal-sized individuals produce less H_2S leading to a less
249 prominent chemical gradient that is not capable of initializing concretion growth (Gaines et
250 al., 2012b). Nevertheless, H_2S can still react with iron from the sediments to form pyrite
251 crystals under anoxic conditions (Raiswell et al., 1993; Schiffbauer et al., 2014). The
252 establishment of anoxic conditions at the time of burial, and H_2S production in the Fezouata
253 Shale lead to the precipitation of framboid and small euhedral crystals in fossils as well as
254 fresh pyrite in non-altered sediments (Fig. 6) (Saleh et al., 2020a). In these sediments, C is
255 also associated with pyrite, possibly suggesting that the original mode of preservation in the
256 Fezouata Shale was comparable to that of the Burgess Shale and the Chengjiang Biota
257 comprising both organic material and authigenic minerals (Gaines et al., 2008; Saleh et al.,
258 2020a,c). As no fossil shows complete pyritization and pyrite precipitation remains rare and
259 tissue-selective, other parameters were also likely controlling pyrite precipitation in the
260 Fezouata Shale (Saleh et al., 2020c). For instance, cuticles of many arthropod taxa are
261 preserved without any pyrite crystals. This can result from H_2S limitation considering that this
262 structure is formed of polysaccharides that are not easily degradable (i.e. quantity of decayed
263 organic matter is not enough to reduce SO_4^{2-}) (Gabbott et al., 2004). However, when
264 comparing internal labile tissues to each other, the model based on H_2S limitation cannot
265 explain why some tissues are pyritized, while others decayed and disappeared (meaning they
266 reduced SO_4^{2-}) without pyritizing (Saleh et al., 2020c). The clue probably lies in Fe

267 availability. Maghemite is found associated with pyrite in some analyzed samples under
268 Raman Spectroscopy (Fig. 6) (Saleh et al., 2020c). Maghemite results from the burial of an
269 original mineral called ferrihydrite $[\text{FeO}(\text{OH})]_8[\text{FeO}(\text{H}_2\text{PO}_4)]$ (Mazzetti and Thistlethwaite,
270 2002). Ferrihydrite is a widely distributed biological mineral (Aldred et al., 2009; Hoda et al.,
271 2013; Dunaief et al., 2014). Under anoxic and sulfated conditions, ferrihydrites release high
272 quantities (~ 87%) of reactive Fe (Li et al., 2006) very fast (i.e. half-life of only 2.8 hours)
273 (Canfield et al., 1992). This is 100 times faster than goethite, 270 times faster than hematite
274 and 2000 times faster than reactive silicates (Canfield et al., 1992). For this reason, in order to
275 understand the patterns of exceptional preservation in the Fezouata Shale, but also in
276 Lagerstätten such as Chengjiang and the Beecher's Trilobite Bed in which pyrite played a role
277 in preserving decay-prone anatomies, three parameters should be taken into account: Fe in
278 sediments, Fe in labile tissues, and H_2S production (Saleh et al., 2020c). Accounting for both
279 pre-burial and anoxic decay, different scenarios emerge and are summarized in Fig. 6.

- 280 • In the first scenario, pre-burial decay is not restricted by any mineralogical phase and
281 burial allowing the establishment of anoxic conditions is delayed. Pyritization does not
282 occur rapidly enough, leaving only body walls (e.g. trilobite carapaces) preserved (Fig.
283 6A).
- 284 • In the second scenario, burial occurs establishing anoxic conditions for pyritization. Fe in
285 burial material is highly reactive leading to the complete pyritization of the organism, if it
286 is buried alive (Fig. 6B). If the animal decayed on the seafloor but the activity of its
287 degradation was controlled by clay/chlorite minerals, the reactivity of Fe from sediments
288 ensures the pyritization of all tissues that survived pre-burial decay (Fig. 6C). Due to
289 extreme iron reactivity, even carapaces of numerous arthropod taxa that provide small
290 quantities of H_2S are found pyritized in sites such as the Beecher's Trilobite Bed (Briggs
291 et al., 1991; Raiswell et al., 1993, 2008).

- 292 • In the third scenario, pre-burial decay is controlled by clay/chlorite minerals. However,
293 after burial, a delay in iron availability in the sediment allows the disappearance of tissues
294 owing to anaerobic decay. The least labile internal tissues will potentially survive anoxic
295 decay and get pyritized once iron from sediments becomes available (Fig. 6D). This
296 scenario explains the selective preservation of guts, while more labile tissues (e.g. nervous
297 systems) are absent (Gutiérrez-Marco et al., 2017).
- 298 • In the fourth scenario, pre-burial decay does not occur at all as if animals were buried
299 alive. However, Fe in this scenario is not reactive (even if it is abundant). Thus, only
300 tissues that are rich originally in iron will get preserved and pyritized, even if they are the
301 most labile ones (Fig. 6E). This scenario can explain the preservation of extremely decay-
302 prone structures such as nervous tissues as pyrite replicates in fossils from the Chengjiang
303 Biota (Ma et al., 2012, 2015; Tanaka et al., 2013; Cong et al., 2014).

304 Considering the decay stages of animals from the Fezouata Shale, the lower availability of
305 iron in sediments in this formation in comparison to sites such as Beecher's Trilobite Bed, and
306 the absence of preserved nervous systems comparable to those occurring in Chengjiang, it is
307 most probable that the taphonomic scenario of most exceptionally preserved fossils found in
308 the Fezouata Shale followed the third scenario (Fig. 6D). However, this pathway is not
309 exclusive, and other scenarios may have accounted for the discovery of only biomineralized
310 parts in some levels from this formation (Fig. 6A). Furthermore, there might have been other
311 parameters facilitating exceptional preservation in the Fezouata Shale. The dominance of
312 storm-induced deposits, the presence of a favorable clay mineralogy and the selective
313 authigenic mineralization surely helped exceptional preservation to occur in this formation.
314 However, these conditions can be acquired in a wide range of marine settings while
315 Ordovician exceptionally preserved biotas remain scarce. The position of the Fezouata Shale

316 near the South pole in a generally cold water environment might have favored the occurrence
317 of exceptional fossil preservation by slowing down decay rates.

318

319 **5. Question 4: post-diagenesis – minimal maturation and extensive modern weathering**

320 Raman spectroscopy-based models for thermal maturation on carbon indicate that the
321 Fezouata Shale sediments were buried at maximum temperatures of 200°C (Rahl et al., 2005;
322 Kouketsu et al., 2014; Saleh et al., 2019, 2020a). Thus, metamorphism *sensu stricto* and
323 organic remain volatilization (Fig. 2) did not occur (Saleh et al., 2019, 2020a). In the absence
324 of metamorphism, calcium leaching and the enrichment of manganese in fossils (e.g. Ca
325 leaching from the skeletal elements of solutan echinoderms and the Mn enrichment around
326 the appendages of marrellomorph arthropods (Fig. 7) in the Fezouata Shale are most probably
327 due to modern weathering (Fig. 2). In the Draa Valley, this formation is exposed to abundant
328 water circulation (Warner et al., 2013). Outcrops are surrounded by abandoned terraces and
329 by numerous water wells (Warner et al., 2013). Circulating waters in arid environments with
330 occasional rain falls similar to the Draa Valley are rich in manganese oxides that can start the
331 oxidation of the tissues that were selectively and three dimensionally preserved in pyrite
332 during early diagenesis in addition to the rest of the fossil (Fig. 7A) (Potter and Rossman,
333 1979; Warner et al., 2013). The resulting products of this reaction are Fe-oxides in the shape
334 of the original pyrite crystals (i.e. euhedral or framboidal) and Mn-sulfates (Fig. 7B) (Larsen
335 and Postma, 1997). If the quantities of Mn are not sufficient to fully oxidize pyrite, pyrite
336 oxidation can further continue through the usage of dissolved oxygen in pore waters (Fig. 7B)
337 and unleash considerable amounts of sulfuric acid (Fig. 7C), thus reducing the pH of the
338 environment and contributing to the dissolution of nearby carbonates (Larsen and Postma,
339 1997) such as the skeletal elements of echinoderms (Fig. 7D). When extensive weathering
340 occurs by circulating waters that are Fe-rich, C can be completely leached from flattened

341 fossils and star-shaped iron oxides can be deposited (Fig. 7E) (Saleh et al., 2020a). These star-
342 shaped and modern iron oxides do not result from pyrite oxidation (Fig. 7) (Saleh et al.,
343 2020a) and they account for the red/orange color of many flattened fossils found in the
344 Fezouata Shale (Fig. 1). Although these minerals are weathering products, they lead to an
345 easier differentiation of biological features from the surrounding sediments. This can be
346 evidenced when comparing extensively weathered and less altered fossils belonging to the
347 same animal group such as marrellomorphs (Fig. 7).

348

349 **6. Question 5: preservation fidelity – underestimation of completely cellular organisms**

350 The Fezouata Shale is major step forward to complete our understanding of the taphonomic
351 mechanisms behind BST preservation in addition to deciphering how these mechanisms
352 influence the patterns of fossil preservation during the Palaeozoic. Burgess Shale-type
353 animals in the Fezouata Biota are preserved just below the SWB. This contrasts with their
354 preservation in much deeper settings during the Cambrian (e.g. the Burgess Shale, Qingjiang
355 Biota) (Gaines, 2014; Fu et al., 2019), and indicates that BST preservation can occur under
356 different bathymetries. Furthermore, the Fezouata Shale shows that BST preservation can
357 occur even if carcasses were exposed on the seafloor prior to their burial as some clay
358 minerals with antibacterial properties can reduce the impact of oxic decay (McMahon et al.,
359 2016; Anderson et al., 2018, 2020a; Saleh et al., 2019). However, this facilitating condition is
360 not enough on its own for soft-structures to preserve because these clay minerals do not stop
361 aerobic bacterial decay; they simply slow it down (McMahon et al., 2016). Most importantly,
362 even if a certain tissue survived oxic degradation it can still decay after burial when different
363 redox conditions and new bacterial communities take over the degradation process (Fig. 6D,
364 E) (Hancy and Antcliffe, 2020). This post-burial degradation can be mitigated by authigenic
365 mineralization such as pyrite or phosphate precipitation (Saleh et al., 2020c; Gueriau et al.,

366 2020) or even by the association of clay minerals such as kaolinite to certain labile anatomies
367 (Anderson et al. 2020b). Authigenic pyritization happens when iron is available and anoxic
368 conditions are established in sediments, but oxidants from the water column (i.e. sulfates) can
369 still reach decaying carcasses (Raiswell et al., 1993, Saleh et al., 2020c). Carbonate cement,
370 found in many Cambrian Lagerstätten, capping event deposits and blocking oxidants from
371 attaining decaying carcasses, are absent from all sediments in the Fezouata Shale which can
372 explain why pyritization is more abundant in this site than in the Walcott Quarry (Burgess
373 Shale) for example (Saleh et al., 2020a). The last main difference between the Fezouata Shale
374 and most Cambrian Lagerstätten is that fossil transport was not operational at the Ordovician
375 site (Martin et al., 2016a; Vaucher et al., 2016, 2017; Saleh et al., 2018, 2020a, b). All these
376 aforementioned preservational discrepancies indicate that there is no single condition that can
377 be considered as a pre-requisite for BST preservation. BST preservation is a trade-off between
378 decay and mineralization (Fig. 6). This trade-off is controlled by interacting parameters (e.g.
379 transport, oxygen, carbonate cement, specific clay minerals, bacteria). This interaction
380 dictates what and how tissues get preserved in a certain site and the impact of this interaction
381 on preservation can be easily investigated by comparing the co-occurrence of biological
382 structures (Fig. 8A) in these deposits, because all animals are originally made of the same
383 types of structures (e.g. mineralized, cuticularized, cellular, sclerotized) (Saleh et al., 2020d).
384 It has been recently evidenced that biological structure association in the Fezouata Shale is
385 significantly different from that at the Burgess Shale and the Chengjiang Biota (Saleh et al.,
386 2020d). Most taxa in the Walcott Quarry and the Chengjiang Biota preserved two to three
387 types of biological structures at the same time (e.g. cuticle and sclerites; or cuticle, sclerites,
388 and cellular tissues, Fig. 8B) (Saleh et al., 2020d). Most taxa in the Fezouata Shale preserved
389 one type of biological structures at a time (Fig. 8B) (Saleh et al., 2020d). Most importantly,
390 the Fezouata Shale systematically failed to preserve soft cellular structures that are in direct

391 contact with seawater in addition to completely cellular organisms. These results suggest that
392 biodiversity is very likely underestimated in the Fezouata Biota (Fig. 8C) (Saleh et al.,
393 2020d). The absence of these structures can be either linked to (1) the fact that cellular
394 structures in direct contact with sea water (e.g. tentacles of hyoliths) are generally the most
395 labile and the fastest to decay (even when clay minerals with antibacterial properties are
396 present) and/or (2) the lack of required conditions for their mineralization (Saleh et al.,
397 2020d). The magnitude of this underestimation can be quantitatively assessed by looking at the
398 proportion of entirely cellular taxa in Cambrian Lagerstätten. In the Walcott Quarry and the
399 Chengjiang Biota, entirely cellular organisms (e.g. chordate, medusoids) constitute 13 and
400 10% respectively of original animal communities in these sites (Saleh et al., 2020d). If the
401 Fezouata Biota had a similar taxonomic composition to the Walcott Quarry and the
402 Chengjiang Biota, the total diversity of this site is most probably underestimated by at least
403 10% (entirely cellular organisms being completely absent). Since this taphonomic bias can be
404 identified and accounted for, the in-situ preservation in the Fezouata Shale becomes an asset
405 for sophisticated ecological studies that examine the evolution of community structure during
406 the Palaeozoic at the transition between the Cambrian Explosion and the Ordovician
407 Radiation. Cambrian ecosystems contrast with Ordovician ones (niche vs dispersal
408 assembled) (Na and Kiessling, 2015; Rasmussen et al., 2019; Stigall et al., 2019). The
409 Fezouata Shale is the only exceptionally preserved biota with hundreds of fossiliferous levels
410 within two stratigraphic intervals (Tremadocian and Floian in age), providing a database that
411 can be used to describe the continuum of ecological change between major Palaeozoic
412 evolutionary events and to decipher if the transition from niche to dispersal-built ecosystems
413 was abrupt or gradual (Harper and Servais, 2018; Rasmussen et al., 2019; Servais et al., 2019;
414 Stigall et al., 2019).

415

416 **7. Conclusions**

417 The Fezouata Shale constitutes a unique record of marine life in the Lower Ordovician. It
418 contains two stratigraphic intervals bearing exceptionally preserved animals (Tremadocian
419 and Floian in age), and documenting an unexpected evolutionary melting pot of Cambrian
420 BST-like holdovers co-occurring with typical members of the Ordovician Radiation. These
421 animals lived in a shallow marine environment that is dominated by storms and were
422 exceptionally preserved *in-situ* just below the SWB. Due to the rarity of storm-induced
423 deposits affecting distal settings, most soft parts decayed prior to their burial. This decay
424 activity was slowed down by a specific clay mineralogy. When favorable conditions for
425 pyritization occurred after burial, pyrite precipitated in specific tissues. In the Fezouata Shale,
426 fossils were originally preserved as carbonaceous compressions with accessory authigenic
427 mineralization in a similar way to Cambrian BST deposits. However, modern water
428 circulation within outcrops leached most of the organic carbon from fossils and oxidized
429 pyrite. Even though the mode of preservation appears to be universal between both Cambrian
430 and Ordovician BST deposits, our review highlights that there are some discrepancies in the
431 mechanisms that lead to this type of preservation. The pre-burial decay in the Fezouata Shale
432 largely contributed to the non-preservation of both soft cellular structures in direct contact
433 with seawater and cellular organisms (e.g. chordates, jellyfish), thus leading to an
434 underestimation of the original diversity of this biota. In spite of this underestimation, the
435 Fezouata Shale is unique in providing an *in-situ*, fully marine community of Burgess Shale-
436 type. The Fezouata Biota represents a series of highly diverse high-latitude communities, at
437 the transition between the Cambrian Explosion and the Ordovician Radiation and as such sits
438 at a critical crossroads in understanding the early evolution of complex animal communities.

439

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753

754 **Figure captions**

755 **Figure 1.** Fossils from the Fezouata Biota. (A) The distribution of classes among fossiliferous
756 levels from the Fezouata Shale. (B) Xiphosuran chelicerate indet. AA.TER.OI.3, (C) hyolith
757 with preserved internal parts YPM515750, (D) *Bavarilla* sp., trilobite with preserved antennae
758 AA.BIZ15.OI.16, (E) *Furca*, marrellomorph arthropod AA.BIZ31.OI.39, (F) *Palaeoscolex?*
759 *tenensis*, palaeoscolecoid worm, AA.BGF2.OI.1, (G) Aglaspidid arthropod, AA-TER-OI-5,
760 (H) *Pirania auraeum*, demosponge AA.JBZ.OI.115, (I) *Thelxiopse*-like arthropod YPM
761 226544, (J) *Tariccoia tazagurtensis*, liwiid arthropod, MGL102155a, (K) *Bohemiaecystis* sp.,
762 stylophoran echinoderm with preserved soft tissues including the water vascular system
763 AA.BIZ15.OI.259, (L) frontal appendages of *Aegirocassis benmoulae*, radiodont arthropod
764 YPM 527123. AA: University Cadi-Ayyad, Marrakesh, YPM: Yale Peabody Museum, MGL:
765 Lausanne University.

766 **Figure 2.** Processes and pathways involved during the transfer of organic matter from the
767 biosphere into the lithosphere (i.e. fossilization). These processes determine which characters
768 from the original morphology are lost or retained (Sansom et al., 2010; Bath Enright, 2018;
769 Parry et al., 2018; Purnell et al., 2018).

770 **Figure 3.** Ordovician lithostratigraphic sequencing of the Zagora area (modified from
771 Marante, 2008) completed with graptolite biostratigraphy for the Fezouata Shale (based on
772 Gutiérrez-Marco and Martin, 2016). Exceptionally preserved fossils occur discontinuously
773 (every 100,000 years) within eccentricity-controlled levels belonging to two stratigraphic
774 intervals (Saleh et al., 2019).

775 **Figure 4.** Sedimentary structures in the Fezouata Shale, typical of a storm-wave dominated
776 environment. (A) Storm bed surrounded by background sediments, (B) bed surface displaying
777 2D oscillatory ripples of centimetric wavelength, (C) storm beds displaying a normal grading,
778 (D) hummocky cross-stratified sandstone showing a metric wavelength, (E) In shallower
779 settings, a tidal modulation of storm waves occurs and is recorded as a repeated stack of
780 larger (low tide) to smaller (high tide) oscillatory structures within fine-grained sandstone, (F)
781 In distal settings, fossils were decaying on the seafloor prior to their burial by storm-induced
782 deposits. Disarticulated skeletal elements of the rhombiferan echinoderm *Macrocyrtella*
783 *bohemica* in green, trilobites in blue, bioturbations in pink, and undetermined elements in
784 beige, (G) most exceptionally preserved BST fossils are preserved *in-situ* under storm-
785 induced beds. Some shelly organisms can be preserved within background sediments because
786 they are resistant to decay and they do not require an event deposition (Kouraiss et al., 2019;
787 Vannier et al., 2019; Saleh et al., 2020a).

788 **Figure 5.** Body size variations of epibenthic, shallow endobenthic, and deep endobenthic taxa
789 along the proximal-distal axis of the Fezouata Shale accordingly with differences in burial
790 rates and oxygenation (OMZ= Oxygen Minimum Zone; SWB: Storm Wave Base).

791 **Figure 6.** Different scenarios of decay and mineralization according to a model based on Fe
792 availability in biological tissues, Fe reactivity in the sediments and H₂S production (Saleh et
793 al., 2020c). (A) Absence of pyrite precipitation due to a delay in burial (*Platypeltoides*
794 *magrebiensis* trilobite; AA.TGR0a.OI.132). (B, C) Most of the organism is pyritized
795 including the body wall (*Triarthrus eatoni* trilobite; YPM.516160) (Farrell et al., 2011). (D)
796 Pyritized parts of the digestive system (*Megistaspis hammondi* trilobite; MGM.6755X)
797 (Gutiérrez-Marco et al., 2017). (E) Pyritized nervous system preserved as a dark brown/black
798 imprint (*Fuxanhuia protensa* arthropod; YKLP.15006) (Ma et al., 2015). The purple spectrum
799 is indicative of chamosite/berthierine identified using X-Ray Diffraction. The red arrows and
800 boxes are examples of pyritized areas identified using Raman Spectroscopy and
801 Backscattered Electronic Microscopy on fossils and fresh sediments. Orange boxes are
802 examples of carbonaceous preservation identified under Raman Spectroscopy and
803 Backscattered Electronic Microscopy as well. MGM: Museo Geominero, Madrid, YKLP:
804 Yunnan Key Laboratory for Paleobiology.

805 **Figure 7.** Modern weathering effect on fossils from the Fezouata Shale. Manganese oxides
806 start the oxidation process (A) followed by dissolved oxygen in pore waters (B). The resulting
807 products of these reactions are respectively manganese sulfates (B) and sulfuric acid (C).
808 Sulfuric acid leaches calcium from the skeletal elements of the fossils (D). If extensive
809 weathering occurs, all C is leached from fossils and replaced by modern, star shaped iron
810 oxides from circulating water (E). This process explains the absence of Ca in the skeleton of a
811 solutan echinoderm (CASG72938) and the enrichment in Mn- deposits surrounding the
812 extensively weathered marrellomorph (AA.BIZ31.OI.39). Modern star-shaped iron oxides
813 that are deposited during this process gave the extensively weathered marrellomorph a
814 red/orange color in comparison to the less weathered marrellomorph (AA.BIZ15.OI.364).
815 CASG: California Academy of Sciences.

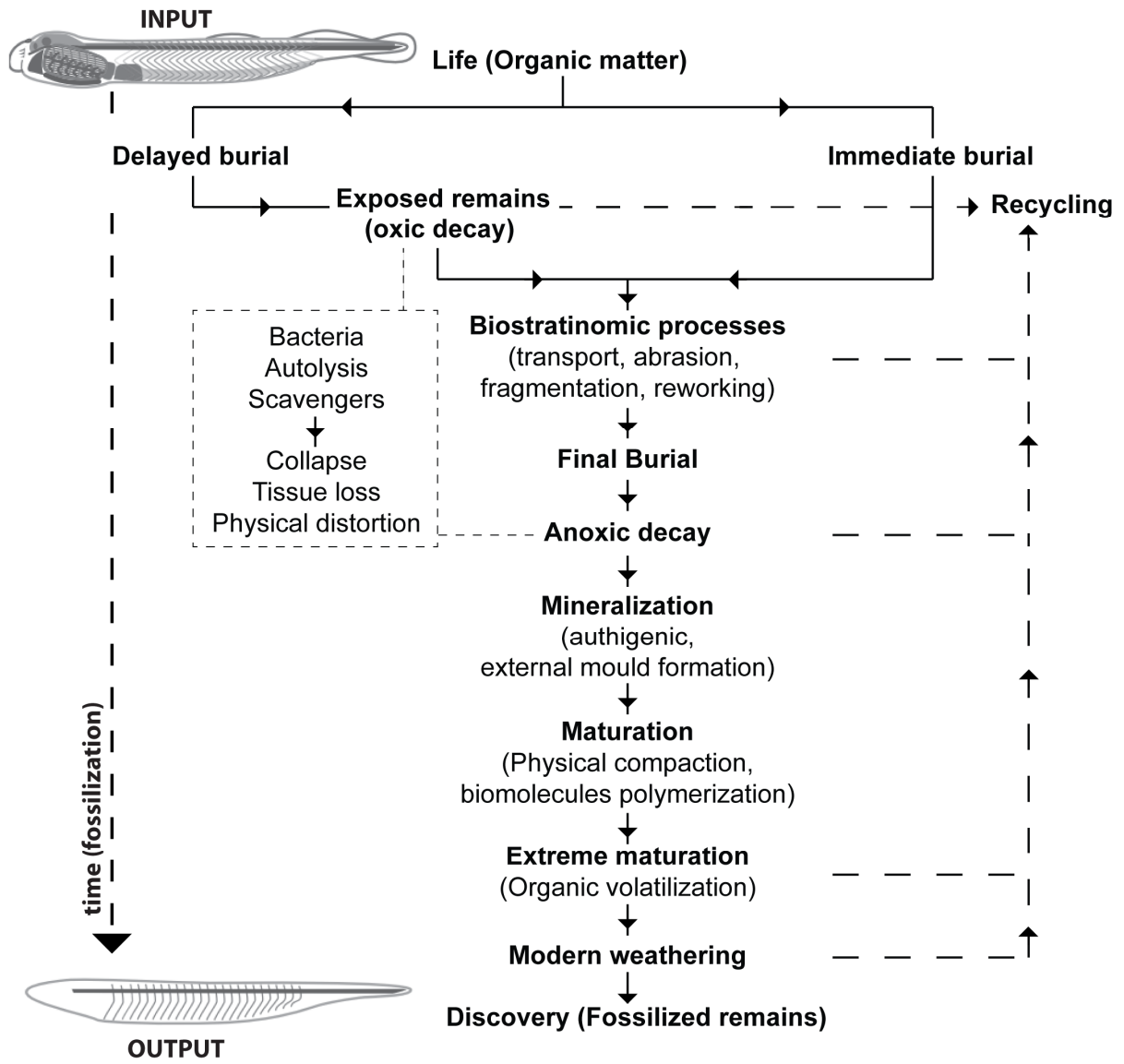
816 **Figure 8.** (A) *Branchiocaris pretiosa*, a crustacean arthropod from the Burgess Shale
817 USNM189028nc preserving three types of biological structures. (B) Differences in the
818 preservational pattern between the Fezouata Shale and the Burgess Shale and the Chengjiang
819 Biota (Saleh et al., 2020d). (C) Exceptionally preserved biotas during the Cambrian and
820 Ordovician plotted on the global diversity curve from Rasmussen et al. (2019). Note that the
821 Fezouata Shale does not preserve completely cellular organisms such as cambroernids and
822 chordates; Cambrian Explosion (CE), Ordovician Radiation (OR). USNM: United States
823 National Museum of Natural history.



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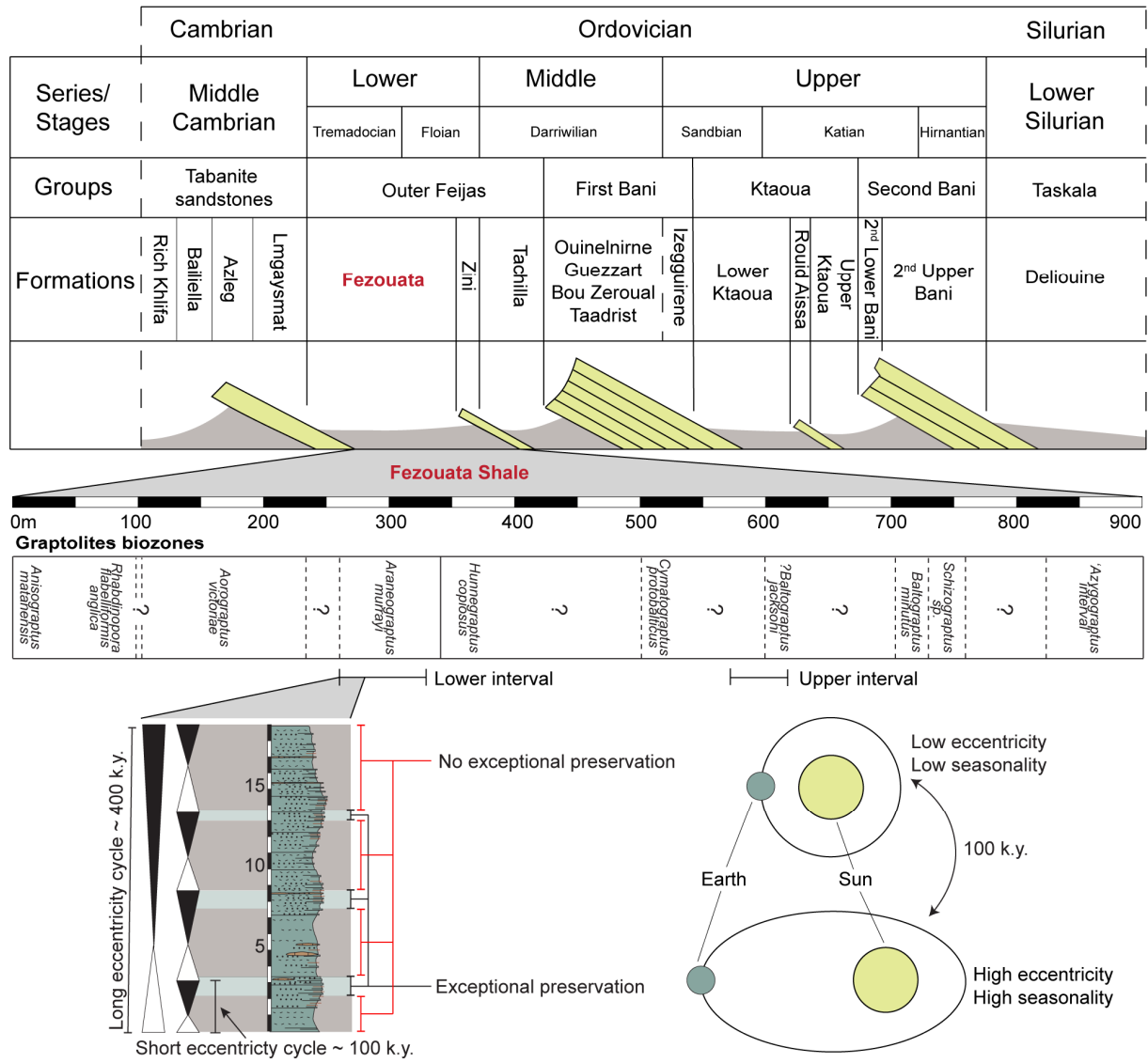
Figure 1



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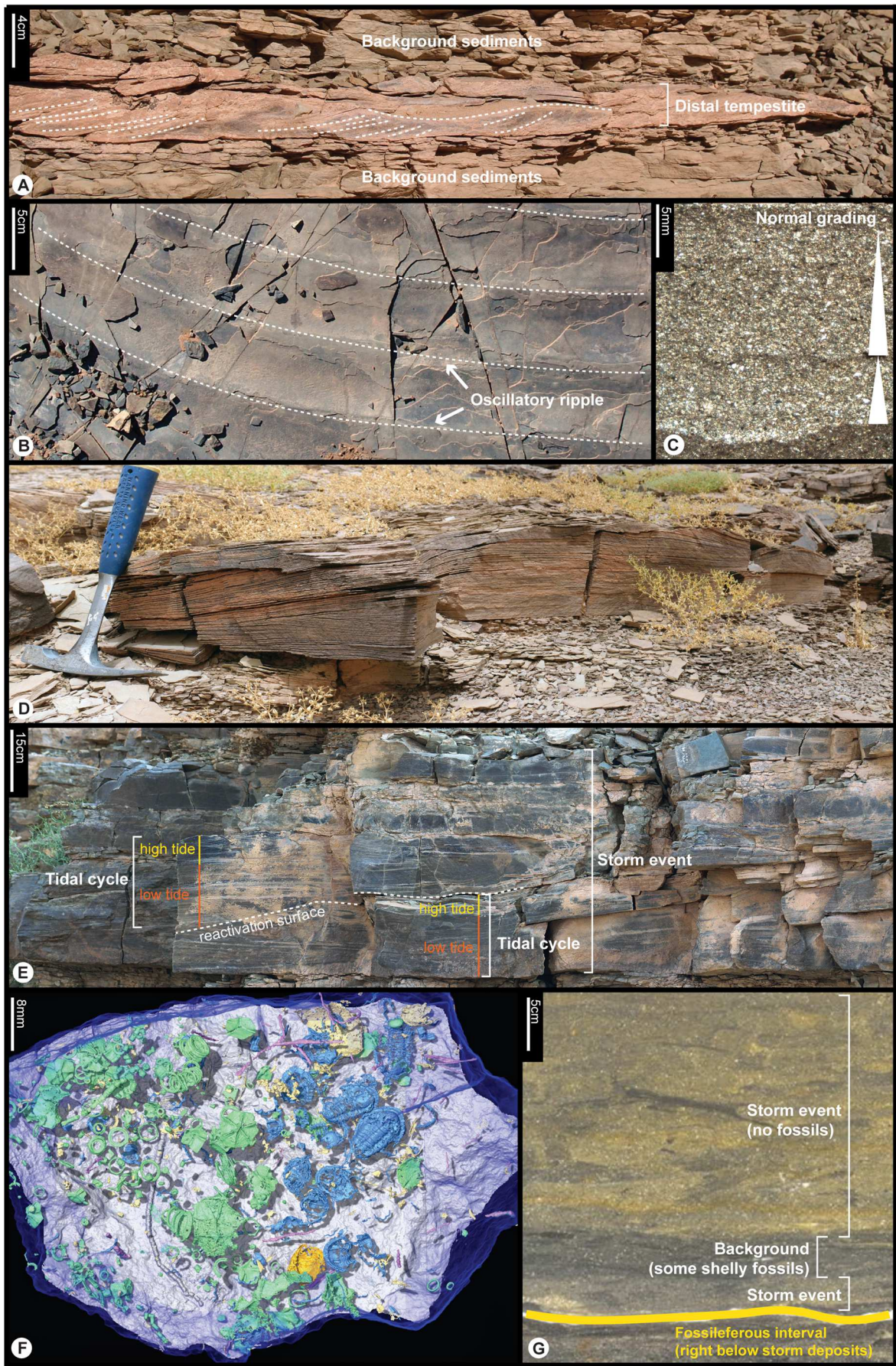
Figure 2



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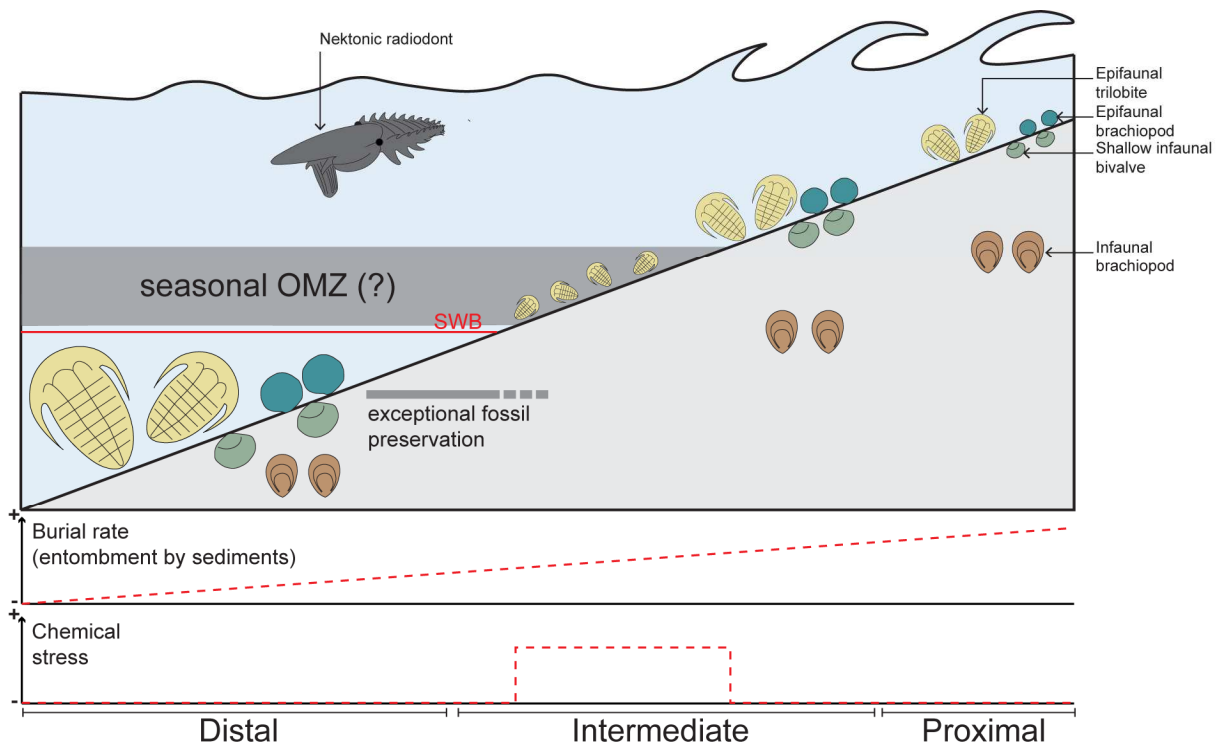
Figure 3



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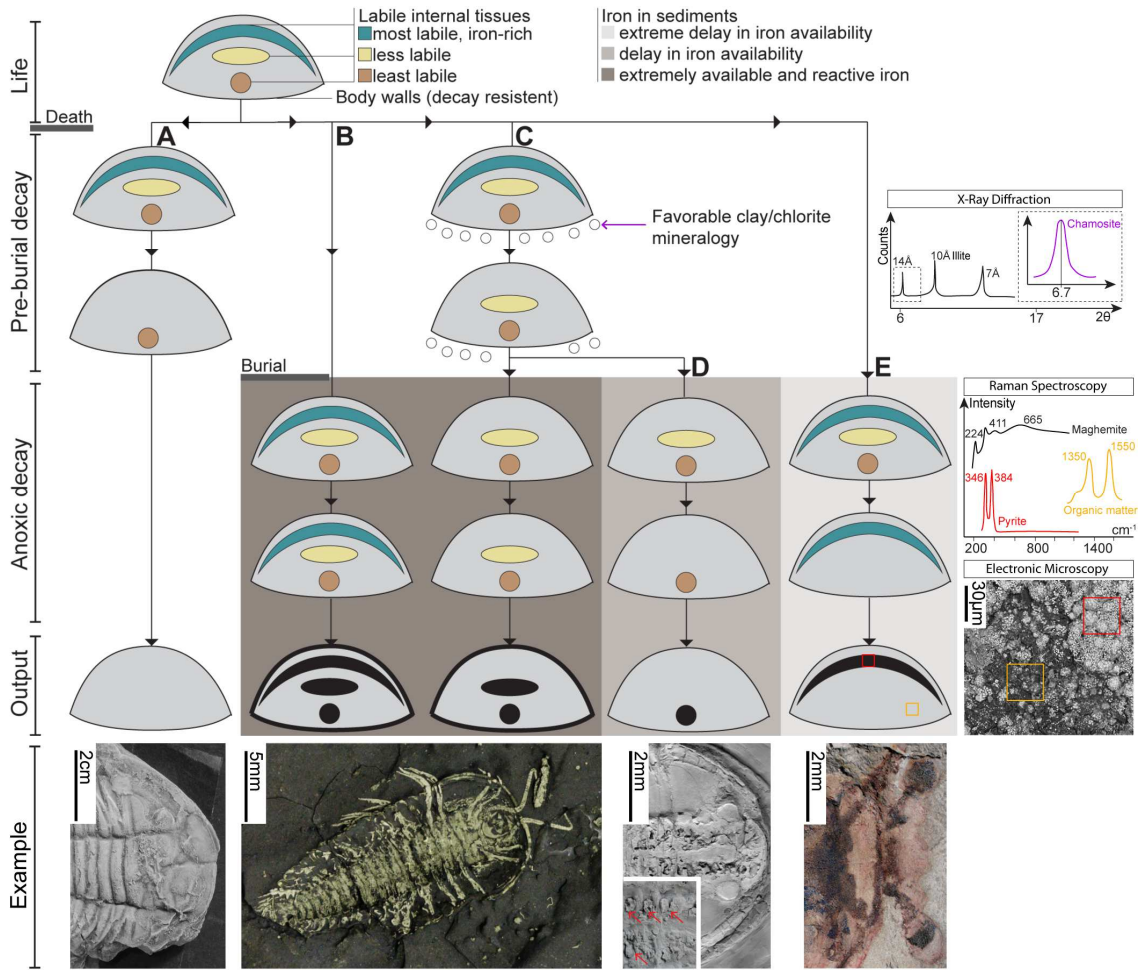
Figure 4



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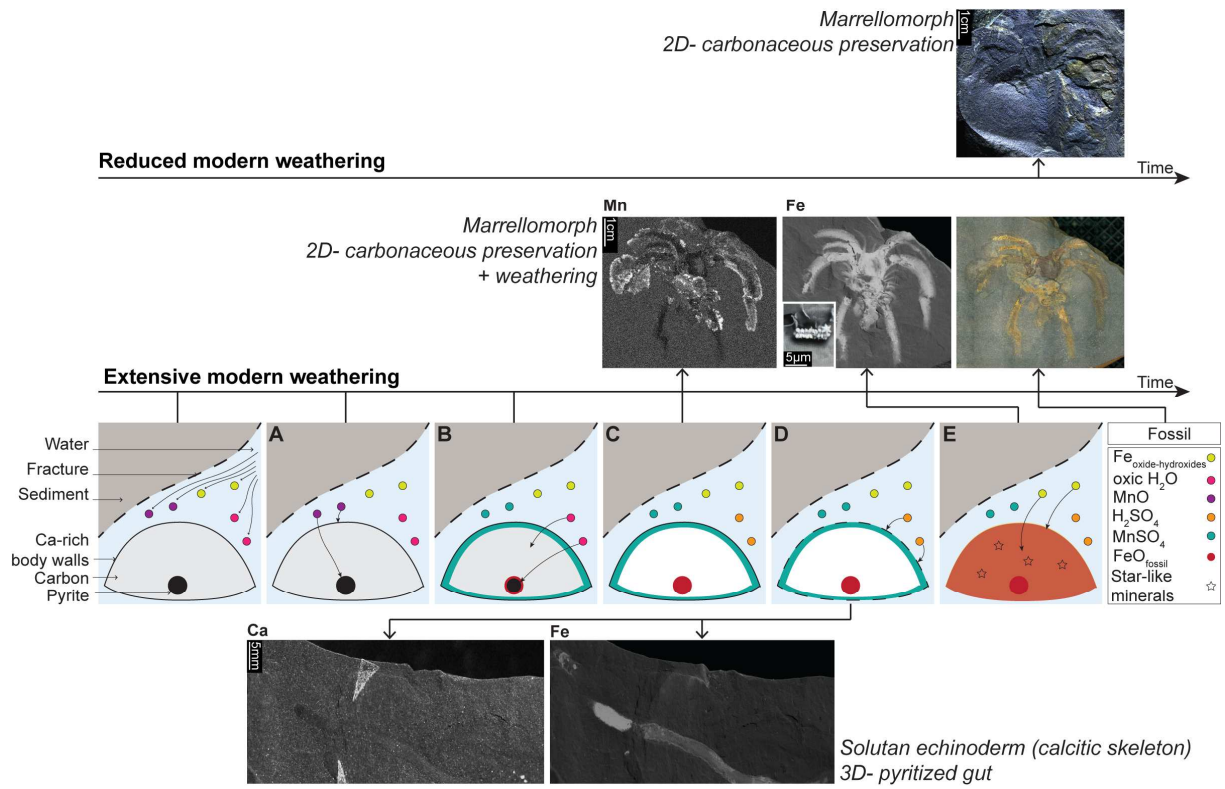
Figure 5



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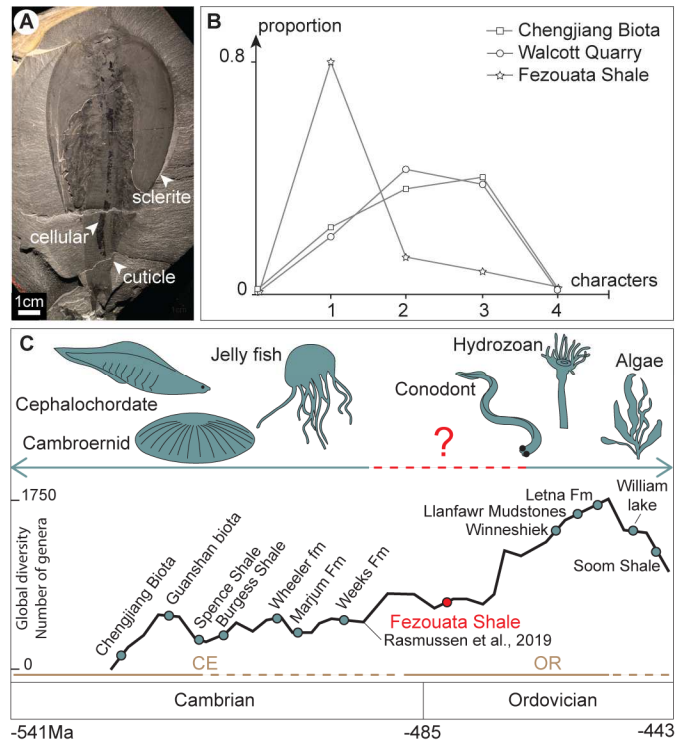
Figure 6



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Figure 7



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Figure 8