

The early Miocene lake of Foieta la Sarra-A in eastern Iberian Peninsula and its relevance for the reconstruction of the Ribesalbes–Alcora Basin palaeoecology

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The Ribesalbes–Alcora Basin (Castelló Province, Spain) contains two lower Miocene units that are rich in fossils. The Unit B contains oil-shale and laminated bituminous dolomitic related to a palaeolake, whereas the Unit C is composed of sandstone and mudstone beds from distal deltaic and shallow lacustrine environments. The La Rinconada and San Chils localities from the Unit B have yielded a fossil assemblage of plants, molluscs, arthropods, and vertebrates, while the localities from the Unit C in the Campisano ravine (Araia/Mas d’Antolino outcrop) are rich in mammalian record. Here we study a new palaeolake deposit of laminated lacustrine limestone beds in the Unit C named Foieta la Sarra-A. This new locality has provided an assemblage of charophytes, terrestrial plants, molluscs, arthropods, and teleosteans. The latter represent the only known fish record from the Ribesalbes–Alcora Basin to date. Although the specimens are generally poorly preserved, the presence of soft-body preservation due to the action of microbial mats at the lake bottom allows considering the Foieta la Sarra-A locality as a Konservat-Lagerstätte. The Foieta la Sarra-A palaeolake had a different water chemistry compared to that represented in the Unit B. Its depth was about a few metres and the water level suffered periodic fluctuations. This new locality sheds light on the palaeoenvironmental dynamics of the Ribesalbes–Alcora Basin during the early Miocene and provides a new approach to the palaeoecological reconstruction of the basin.

Key words: Characeae, Poales, Gastropoda, Cladocera, Insecta, Teleostei, palaeoenvironment, taphonomy, palaeoecology, lacustrine basin, Neogene, Konservat-Lagerstätte, Spain.

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Introduction

Lakes are remarkable ecological archives due to their high sensitivity to climatic changes, a characteristic that can also be used in deep time (Cohen 2003). Indeed, the fossil content and sedimentary record of palaeolakes provide valuable information, such as palaeoclimatic, chemical, and temperature conditions (Talbot and Allen 1996). During the Miocene, a large portion of the European continent was covered by lakes (e.g., Neubauer et al. 2015a, b; Mandic et al. 2019; Vasilyan 2020). Those which developed in the Iberian Peninsula produced exceptionally well-preserved fossil assemblages with Konservat-Lagerstätte characteristics (e.g., McNamara et al. 2012; Peñalver et al. 2016). Remarkable Iberian Konservat-Lagerstätten in Miocene lacustrine deposits are the fossil localities of Rubielos de Mora (Teruel Province), Bicorn (Valencia Province), La Rinconada and San Chils (the latter two in Castelló Province). These well-preserved and diverse fossil assemblages originated under anoxic conditions at the lake bottom with presence of microbial mats (Peñalver 2002; Peñalver and Gaudant 2010).

The Ribesalbes–Alcora Basin is a Neogene complex graben covering around 150 km² located in SE Iberian Range (Anadón et al. 1989; Fig. 1A). This basin is well-known for the mining activities leading to oil-shale extraction in the La Rinconada (Ribesalbes) and San Chils (l'Alcora) mines until the early 20th century and clay extraction in the Campisano ravine (Araia/Mas d'Antolino area) which continues nowadays (Peñalver et al. 2016; Crespo 2017). The Campisano ravine contains seven sections with mammalian fossil record that have been proposed as a Site of Geological Interest (Costa-Pérez et al. 2019), i.e., Mas dels Coixos, Mas de Torner, Araia Cantera Sud, Barranc de Campisano, Foieta la Sarra, Mas d'Antolino B and Corral de Brisca (Fig. 1A). La Rinconada is a well-known locality that has provided an outstanding record of plants, molluscs, ostracods, arachnids, insects, amphibians, bird feathers and coprolites (Peñalver et al. 2016). Furthermore, a fossil assemblage from the San Chils locality has been recently described showing a strong similarity to the assemblage of La Rinconada (Álvarez-Parra and Peñalver 2019). Both localities have been interpreted as different areas of the same meromictic palaeolake, mostly contemporaneous, that occupied the basin during the early Miocene (Álvarez-Parra and Peñalver 2019). The exceptional preservation of their fossil record is related to the presence of microbial mats at the lake bottom (Peñalver et al. 2016). Interestingly, these fossil localities lack fish record, which might be related to an endorheic origin of the lake or an inappropriate water chemistry for the species that could reach the lake (Peñalver et al. 2016).

Forty-five fossiliferous levels rich in mammal remains have been recently described by Crespo et al. (2019c), in addition to the ones previously studied by Agustí et al. (1988), in the seven Campisano ravine sections. Their general features, together with the geological and preliminary biostratigraphic framework, were published by Crespo et al. (2019c), who de-

scribed the geology of the area and placed the sections under study a stratigraphic context. In addition, many of the small mammal groups found in these sections have already been studied, including two new species of bat and dimyid (Furió et al. 2012; Crespo 2017; Crespo et al. 2019a, b, 2020a, b, 2021, in press). Among the different studied sections from these localities, the Foieta la Sarra section includes the Foieta la Sarra-1 locality, which yields a rich record of vertebrate remains (Crespo et al. 2019c). The faunal assemblage of the Foieta la Sarra-1 locality listed by Crespo et al. (2019c) is: *Amphiperatherium frequens erkertshofense* (Koenigswald, 1970), *Heteroxerus rubricati* Crusafont, Villalta, and Truyols, 1955, *Megacricetodon primitivus* (Freudenthal, 1963), *Democricetodon decipiens* (Freudenthal and Daams, 1988), *Eumyarion weinfurteri* (Schaub and Zapfe, 1953), *Microdyromys koenigswaldi* De Bruijn, 1966, *Peridyromys murinus* (Pomel, 1853), *Pseudodryomys ibericus* De Bruijn, 1966, *Simplomys julii* (Daams, 1989), *Glirudinus undosus* Mayr, 1979, *Ligerimys ellipticus* Daams, 1976, *Galerix symeonidisi* Doukas, 1986, *Talpidae* sp. indet., and cf. *Soricella discrepans* Doben-Florin, 1964. The preliminary study of the palaeoecology of this mammalian assemblage by Crespo (2017) indicated an increase of the open forest, as well as higher humidity and temperature than the oldest localities from the Campisano ravine.

Here, we describe the fossil record and the palaeoecology of a new lower Miocene locality we name Foieta la Sarra-A, which namely consists of laminated lacustrine limestone beds located at the base of the Foieta la Sarra section. Our main goal is to infer the taphonomic and palaeoecological conditions of this outcrop. Together with the current data from La Rinconada and San Chils localities, our new findings shed light on the palaeoenvironmental dynamics of the Ribesalbes–Alcora Basin during the early Miocene.

Abbreviations.—EDX, X-ray energy microanalysis.

Geological setting

The deposits of the Ribesalbes–Alcora Basin are composed of detrital and carbonatic materials deposited in alluvial and lacustrine environments overlying a Mesozoic basement (Anadón et al. 1989) and divided into the Ribesalbes sequence (early to middle Miocene in age) and the Alcora sequence (late Miocene in age). The Ribesalbes sequence includes five depositional units (A to E from the base to the top) defined by Anadón et al. (1989). The Unit A consists of a 300 m interval of breccia of Mesozoic limestone clasts with minor interbedded red sandstone and mudstone. The Unit B is characterised by oil-shales and laminated bituminous dolomicrite of the La Rinconada and San Chils localities, which yield a rich and diverse lacustrine fossil record. The Unit C crops out in the Campisano ravine, where the mammalian fossiliferous sites are located. This unit is composed of sandstone beds ranging in thickness between 15

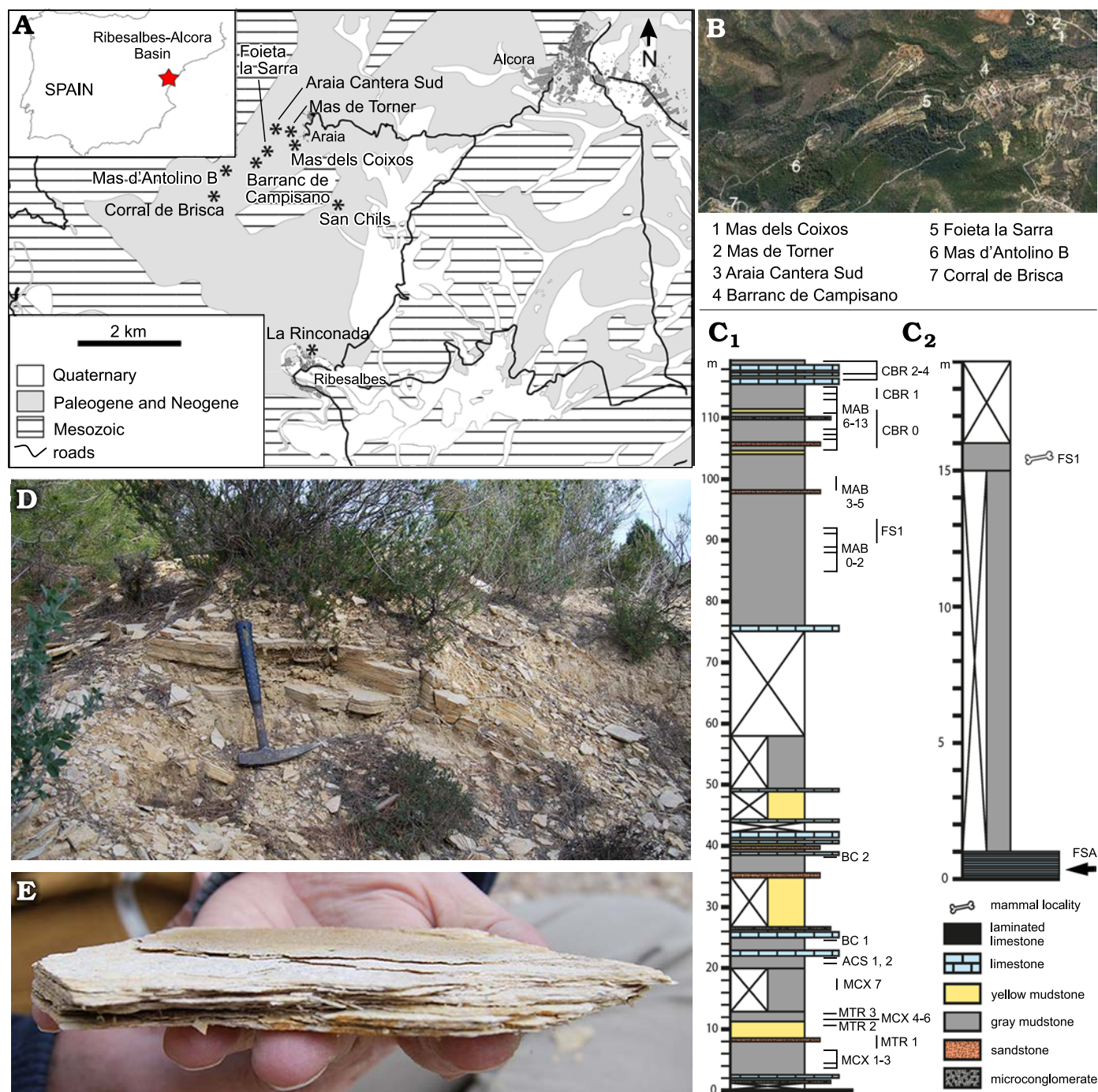


Fig. 1. **A, B.** Geographic and geological location of the Ribesalbes–Alcora Basin (Castelló Province, Valencian Community, Spain) with indication of the Campisano ravine (1–7), San Chils, and La Rinconada localities. **C.** Synthetic stratigraphic column of the Campisano ravine (C₁), and Foieta la Sarra section (C₂), showing the location of the Foieta la Sarra-1 (FS1) and the studied Foieta la Sarra-A (FSA) mammalian fossiliferous localities. **D.** Fossiliferous laminated slabs of laminated lacustrine limestone at the lower Miocene Foieta la Sarra-A locality. **E.** Detail of the fine lamination of the Foieta la Sarra-A of the Foieta la Sarra-A laminated limestone rocks. Abbreviations: ACS, Araia Cantera Sud; CBR, Corral de Brisca; BC, Barranc de Campisano; MAB, Mas d’Antolino B; MCX, Mas dels Coixos; MTR, Mas de Torner. A, B modified from Crespo (2017), Crespo et al. (2019c).

cm and 1.5 m, and which show cross and laminated bedding that alternates with thick massive mudstone strata interbedded with a few dolomitic and calcareous layers. These facies were interpreted by Anadón et al. (1989) as distal deltaic and shallow lacustrine environments. The Unit D represents an olistostrome of Cretaceous rocks, reflecting tectonic activity. Lastly, the Unit E is composed of thin bedded limestone

strata rich in ostracods and charophytes, with interbedded mudstone horizons. On the other hand, the upper Alcora sequence consists of conglomerate beds related to alluvial deposits up to 200 m thick. This sequence displays a continuous transition without unconformities over the Ribesalbes sequence in the Campisano ravine (Anadón et al. 1989).

A synthetic stratigraphic column (Fig. 1C₁) and a geolo-

gical study of the Campisano ravine was presented by Crespo et al. (2019c). The stratigraphic section is 120 m thick and it is chiefly formed by mudstone and limestone beds, including some sandstone and microconglomerate layers. In the Campisano ravine, the Foietta la Sarra section is correlated to the lower part of the Mas d'Antolino B section (Fig. 1C₂). The Foietta la Sarra stratigraphic section is 19 m thick and it consists of grey mudstone. A 1 m thick laminated lacustrine limestone bed, which corresponds to the Foietta la Sarra-A locality, can be distinguished at the base of the Foietta la Sarra section. The mammalian fossil site is located at the top of the Foietta la Sarra section, 16 m above the base, within grey sandy mudstone beds.

According to Crespo et al. (2019b), the presence of the genera *Megacricetodon*, *Ligerimys*, and *Democricetodon* in the Foietta la Sarra-1 locality is correlated with the Biozone C of the MN4 (early Aragonian, early Miocene, 16.49–15.94 Ma), based on the comparison with the Aragonian type area in the Calatayud–Montalbán Basin (García-Paredes et al. 2016). The Biozone C of the MN4 is divided in two sub-biozones, i.e., Ca and Cb, with the limit between them at approx. 16.20 Ma after Van der Meulen et al. (2012). The limit between these sub-biozones is present at the Campisano ravine (Crespo et al. 2019c). Specifically, the Foietta la Sarra-1 locality has provided the oldest record of *Eumyarion weinfurteri* and *Ligerimys ellipticus* within the biostratigraphy of the Campisano ravine localities and in the basin (Crespo et al. 2019c). Considering lower and upper stratigraphic levels, Foietta la Sarra-1 is older than the level Mas d'Antolino B-3 due to the high abundance of *Megacricetodon* and the scarcity of *Democricetodon*, which are probably correlated with the sub-biozone Ca from the Calatayud–Montalbán Basin. Foietta la Sarra-1 is younger than the level Mas d'Antolino B-0B because of the presence of both the genus *Eumyarion* and the species *L. ellipticus*, which are probably correlated with the sub-biozone Cb from the Calatayud–Montalbán Basin (Crespo et al. 2019, in press). Considering this information and the stratigraphic position of Foietta la Sarra-A locality studied here, its age could be approximately 16.20 Ma (early Miocene), and thus located at the base of the Cb sub-biozone.

Material and methods

The material studied herein includes compressed fossils of plants, molluscs, arthropods, and fish recorded in 45 laminated slabs excavated from the Foietta la Sarra-A locality at the Foietta la Sarra section near Araia d'Alcora (l'Alcora, Castelló Province, Spain; Fig. 1). The fossil specimens from Foietta la Sarra-A are housed at the Museu de la Universitat de València de Historia Natural (MUVHN) in Burjassot (Valencian Community, Spain). The sedimentary rock exposed at Foietta la Sarra-A does not show significant signals of weathering, although several Recent small roots and fungal mycelia were removed from the slabs. The rock

slabs were observed with a Leica MS5 stereomicroscope. To enhance visibility of some microfossils, slabs were submerged in ethanol. Specimens were photographed using a Leica DMS1000 stereomicroscope with an attached digital camera, and the drawings were made using a camera lucida attached to the Leica MS5 stereomicroscope. All of these methods took place at the Department of Botany and Geology of the Universitat de València.

The micromorphology of the samples was imaged using a FEI INSPECT (5350 NE Dawson Creek Drive Hillsboro, Oregon 97124, USA) Scanning Electron Microscope (SEM) at the Museo Nacional de Ciencias Naturales (MNCN) in Madrid (Spain). The SEM microscope in low vacuum mode allows hydrated samples to be studied in their original state using the large field detector (LFD), since it is close to the sample in order to avoid electron losses. In addition, the samples were observed with the Backscattering Electron Detector (BSED), which were conductive in high vacuum mode. The SEM resolution at low vacuum was 4.0 nm at 30 kV (BSED); the accelerating voltage was 20 kV, low vacuum 0.50 torr, and 10 mm of working distance. The X-ray energy microanalysis (EDX) of the ephippial microsculpture, chironomid larval galleries and fish remains were conducted with an energy-dispersive X-ray spectrometer (INCA Energy 200 energy dispersive system, Oxford Instruments) at the MNCN (Madrid, Spain).

Anatomic terminology for fish remains follows Lagler (1947), Daniels (1996), Schultze (2015) and Bräger and Moritz (2016).

Foietta la Sarra-A fossil assemblage

The Foietta la Sarra-A locality has yielded a diverse record of plants (including charophytes and embryophytes), molluscs, arthropods (including crustaceans and insects) and vertebrates represented by fish. The botanical and faunal taxa of Foietta la Sarra-A are available in Table 1.

Plants.—Three grey-yellowish limestone intervals containing abundant charophyte remains have been distinguished in the Foietta la Sarra-A (Fig. 2A, B). These intervals are exclusively composed of compressed vegetative (thallus) fragments and poorly preserved gyrogonites (calcified fructifications of Characeae) forming a type of calcareous sedimentary rocks defined as characeite by Soulié-Märsche et al. (2010). The flattened charophyte stems studied herein consist of isostichous corticated thalli. Considering that living charophyte genera have not changed their vegetative structure since the Miocene, these charophyte thalli can be attributed to the genus *Chara* since among the living charophytes only this genus develops corticated stems (Soulié-Märsche et al. 2010). Complete articulated thalli would be required for a species-level attribution. Few poorly preserved gyrogonites occur among charophyte stems (Fig. 2B). The absence of complete gyrogonites and their poor preservation

Table 1. Botanical and faunal record from the lower Miocene Foietta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain).

Charophyta	
Charophyceae	
Charales	
Characeae	<i>Chara</i> sp.
Embryophyta	
Monocotyledoneae	
Poales	Poaceae, Cyperaceae, Sparganiaceae or Juncaceae indet.
Dicotyledoneae	
Incertae sedis	<i>Dicotylophyllum</i> sp.
Malpighiales	
Salicaceae	<i>Salix?</i> sp.
Mollusca	
Gastropoda	
Littorinimorpha	Hydrobiidae? indet.
Hygrophila	Lymnaeidae indet.
Planorbidae	<i>Ferrissia</i> sp. <i>Gyraulus</i> sp.
Arthropoda	
Branchiopoda	Ostracoda indet.
Cladocera	
Daphniidae	<i>Daphnia (Ctenodaphnia)</i> sp.
Insecta	
Coleoptera	Coleoptera indet.
Hemiptera	Pentatomidae? indet.
Diptera	Psychodidae? indet.
Vertebrata	
Actinopterygii	
Teleostei	Teleostei indet.
Cyprinodontiformes	
Cyprinodontidae	cf. <i>Aphanius</i> sp.

hinder a detailed taxonomic attribution. Several molds and casts of planorbid gastropods (*Gyraulus* sp.) also occur associated to these characeite intervals.

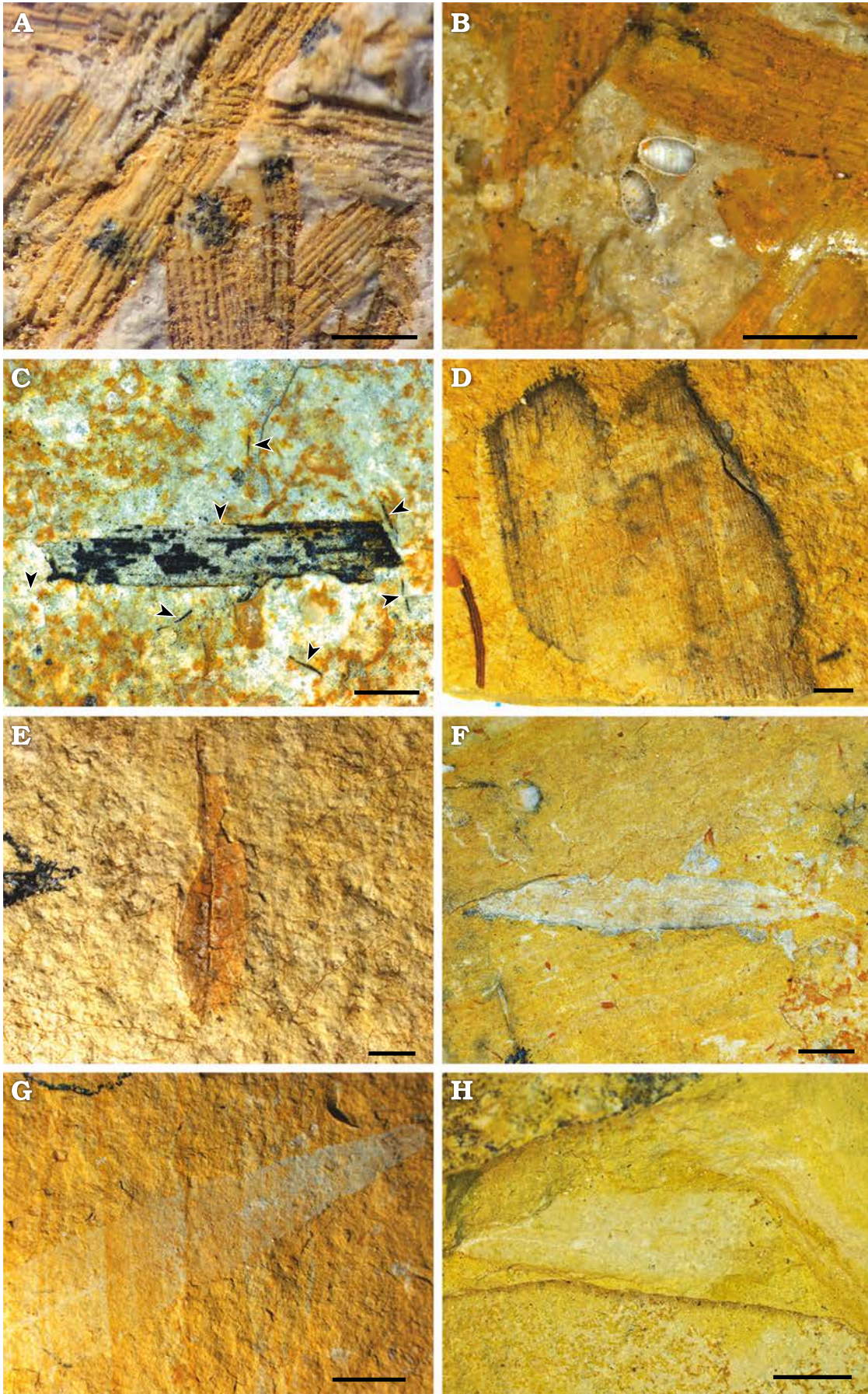
Several microlevels of Foietta la Sarra-A show accumulations of burned plant remains or fusinite (Fig. 2C) as described from La Rinconada and San Chils localities (Peñalver et al. 2016; Álvarez-Parra and Peñalver 2019), possibly resulting from wildfires near the palaeolake. Undetermined seeds with slightly bilobate apex have been found, although their preservation is poor. The Foietta la Sarra-A plant record also includes leaves, but their taxonomic affinities are challenging to determine. A large leaf remain and scarce small fragments with straight margins and parallel veins (Fig. 2D) are similar to specimens of *Typha latissima* A. Braun, 1851 in Heer, 1855 (Typhaceae) found in the Miocene Rubielos de Mora locality (Barrón and Diéguez 2001); this type of plant remain is also found in La Rinconada and San Chils localities. However, they are usually poorly preserved and their morphotype could correspond to Poaceae, Cyperaceae, Sparganiaceae or Juncaceae belonging to the order Poales (Peñalver et al. 2016; Álvarez-Parra and Peñalver 2019). Two

slender leaves with distal areas partly preserved, 11.6 mm long and 2.9 mm maximum wide and 14.3 mm long and 2.12 mm maximum wide, respectively (Fig. 2E, F), seem to be similar to the morphotypes of *Dicotylophyllum* sp. present in La Rinconada (Barrón and Postigo-Mijarra 2011). Two poorly preserved leaf remains with around 6 mm maximum width have been tentatively determined as *Salix?* sp. (Salicaceae) based on the straight, slightly parallel margins and their similarity to the specimens of *Salix lavateri* A. Braun emend. Hantke, 1851 from La Rinconada (Barrón and Postigo-Mijarra 2011; Fig. 2G, H).

Molluscs.—The gastropod assemblage recorded in the studied samples is poorly preserved. Almost all specimens consist of moulds and impressions, which hinder accurate taxonomic attributions. The assemblage consists of at least four freshwater morphotypes, one of which belongs to the superfamily Truncatelloidea and the rest to the superfamily Lymnaeoidea. The Truncatelloidea superfamily is represented in Foietta la Sarra-A, with 15 recognised specimens in seven of the examined samples, by a single morphotype (Fig. 3A–C). Due to the incompleteness of the specimens and their poor preservation, they may belong to Hydrobiidae or Bithyniidae. However, the general ovate-conical morphology of the specimens, with convex spiral whorls and, above all, their small size, suggest the attribution to the family Hydrobiidae.

Regarding the three morphotypes of the superfamily Lymnaeoidea, 120 specimens belonging to the families Lymnaeidae and Planorbidae have been studied. Lymnaeidae is represented by one morphotype, from which eight specimens have been identified (Fig. 3D–G). Seven correspond to fragments of adult specimens (Fig. 3D, E) and one to a juvenile (Fig. 3F, G). The fragments of the adults are very incomplete but show elongate morphology, a last round of rapid growth, and ornamentation of growth constrictions in the vicinity of the aperture. The juvenile specimen, with dimensions of up to 6.7 mm long and 3.7 mm wide, is compressed and deteriorated. Nevertheless, an elongate-ovate shell with three whorls is visible, its spire being short and conical, with a last whorl enlarging rapidly. The assignment at the genus level is challenging, but the characteristics observed match those of the genera *Stagnicola* and *Radix*. Vilanova y Piera (1859) cited the presence of “Lymnaeas” in several areas of the Castelló Province, including the “Balsa de Fanzara” (the classical name for the Cenozoic sediments of the Ribesalbes–Alcora Basin), although this author did not provide further information. In any case, we prefer to keep the determination of the lymnaeid morphotype at the family level.

Two species of the family Planorbidae have been identified in Foietta la Sarra-A, i.e., *Ferrissia* sp. (Fig. 3H–L), with 41 specimens present in 16 of the studied laminated slabs, and *Gyraulus* sp. (Fig. 3M–R), with 71 specimens in 21 of the slabs. The specimens of *Ferrissia*, the largest with 5.90 mm long and 3.89 mm wide, have a limpet-like morphology with an elliptical basal outline. The anterior shell portion is



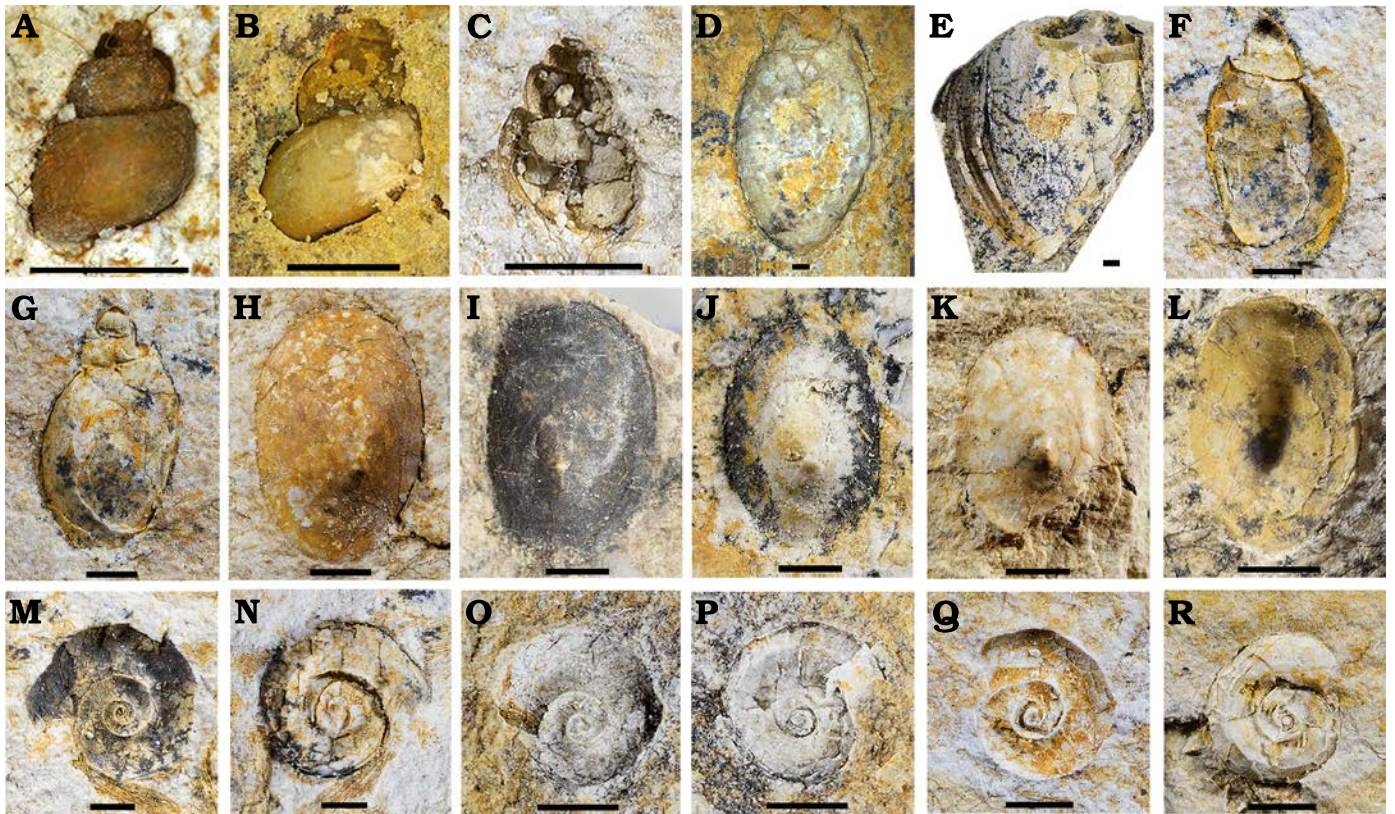


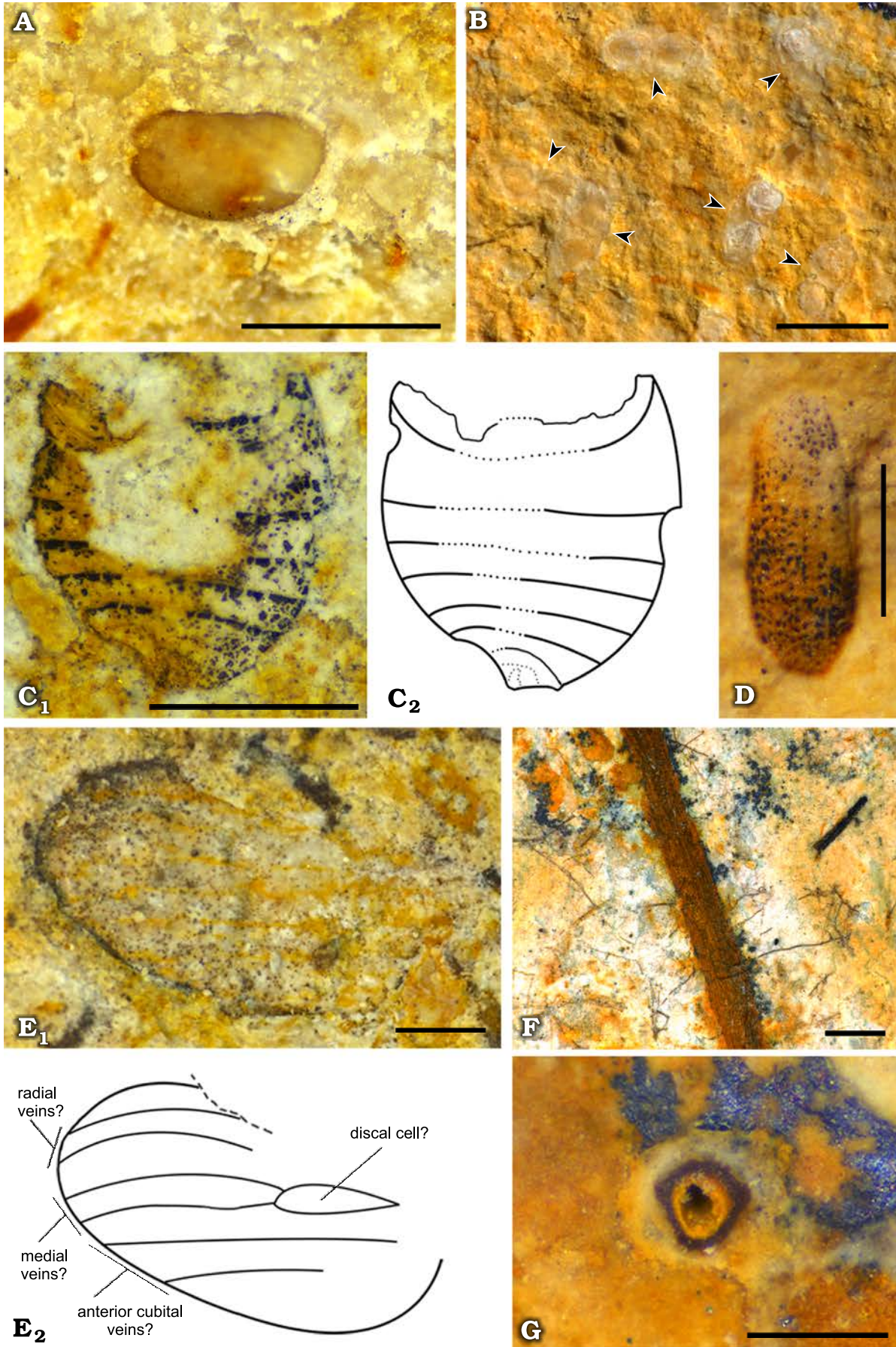
Fig. 3. Molluscs from the lower Miocene Foieta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). A–C. Hydrobiidae? indet. specimens (A, FS-9A-1; B, FS-44C-2; C, FS-15-1). D–G. Lymnaeidae indet. specimens (D, FS-10-1; E, FS-39-1; F, FS-28B-2; G, FS-28A-2). H–L. *Ferrissia* sp. specimens (H, FS-16-1; I, FS-16-3; J, FS-13A-1; K, FS-31C-3; L, FS-39-2). M–R. *Gyraulus* sp. specimens (M, FS-42A-1; N, FS-42B-1; O, FS-35B-1; P, FS-35A-1; Q, FS-7B-1; R, FS-7A-1). Scale bars 1 mm.

slightly broader and convex, the flanks are nearly straight or slightly convex, and the posterior part is convex. The shell apex is slightly deflected to the right, its posterior portion is slightly concave, and its anterior portion is weakly convex in lateral view. The ornamentation is poorly preserved. These specimens of *Ferrissia* from Foieta la Sarra-A locality differ from the middle Miocene *Ferrissia illyrica* (Neumayr, 1880) from the Dinaride Lake System (Neubauer et al. 2011, 2013, 2015c) by its narrower contour. The early–middle Miocene *Ferrissia wittmanni* (Schlickum, 1964) from Southern Germany and Austria (Harzhauser and Kowalke 2002; Kowalke and Reichenbacher 2005; Harzhauser et al. 2014) also has a narrower profile, as well as concave flanks and a cup-shaped apex. Moreover, the late Miocene *Ferrissia truci* Wautier, 1975, from France (Wautier 1975) differs from the Foieta la Sarra-A specimens by its more elongate contour, as well as the elongate apical region with a rounded apex located far back and strongly inclined to the right. The shape of the shell of the studied specimens is particularly similar to that of *Ferrissia depridita* (Desmarest, 1814), a common species in the middle Miocene of Central

Europe. However, the general degree of preservation and particularly the ornamentation prevent us from assigning our specimens to the latter species. On the other hand, *Gyraulus* sp. is the most abundant gastropod species in the studied samples. However, all fossils attributed to this taxon correspond to incomplete moulds and impressions, which hinders the determination to species level. However, the characteristics of this morphotype, whose shell reaches four whorls among the largest specimens, are consistent with those of *Gyraulus* based on the growth pattern (with whorls moderately increasing in diameter), the estimated dimensions among the studied specimens (with a maximum diameter of 5.18 mm), and the ornamentation of growth lines, i.e., prosocline in umbilical view and prosocyrct in apical view.

Arthropods.—Crustaceans and insects represent the hitherto recovered arthropod record from the Foieta la Sarra-A locality. Crustaceans include Ostracoda and Cladocera. Ostracod shells 1–2 mm long, 0.5–1 mm wide without apparent ornamentation have been found (Fig. 4A); they have been recorded both as isolated specimens and as mass records (blooms) in the laminated slabs. The poor preservation

← Fig. 2. Plant remains from the lower Miocene Foieta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). A, B. *Chara* sp. corticated isostichous thalli (FS-3A) and two gyrogonites surrounded by thalli (FS-3B). C. Burned plant remains or fusinite of diverse size (arrowheads) (FS-8A-1). D. Poorly preserved Poaceae, Cyperaceae, Sparganiaceae or Juncaceae leaf remain (FS-21B-1). E, F. *Dicotylophyllum* sp. leaf remains (E, FS-22A-1; F, FS-24B-1). G, H. *Salix*? sp. leaf remains (G, FS-19B-2; H, FS-40-1). Scale bars A–C, 1 mm; D–F, 2 mm; G, H, 4 mm.



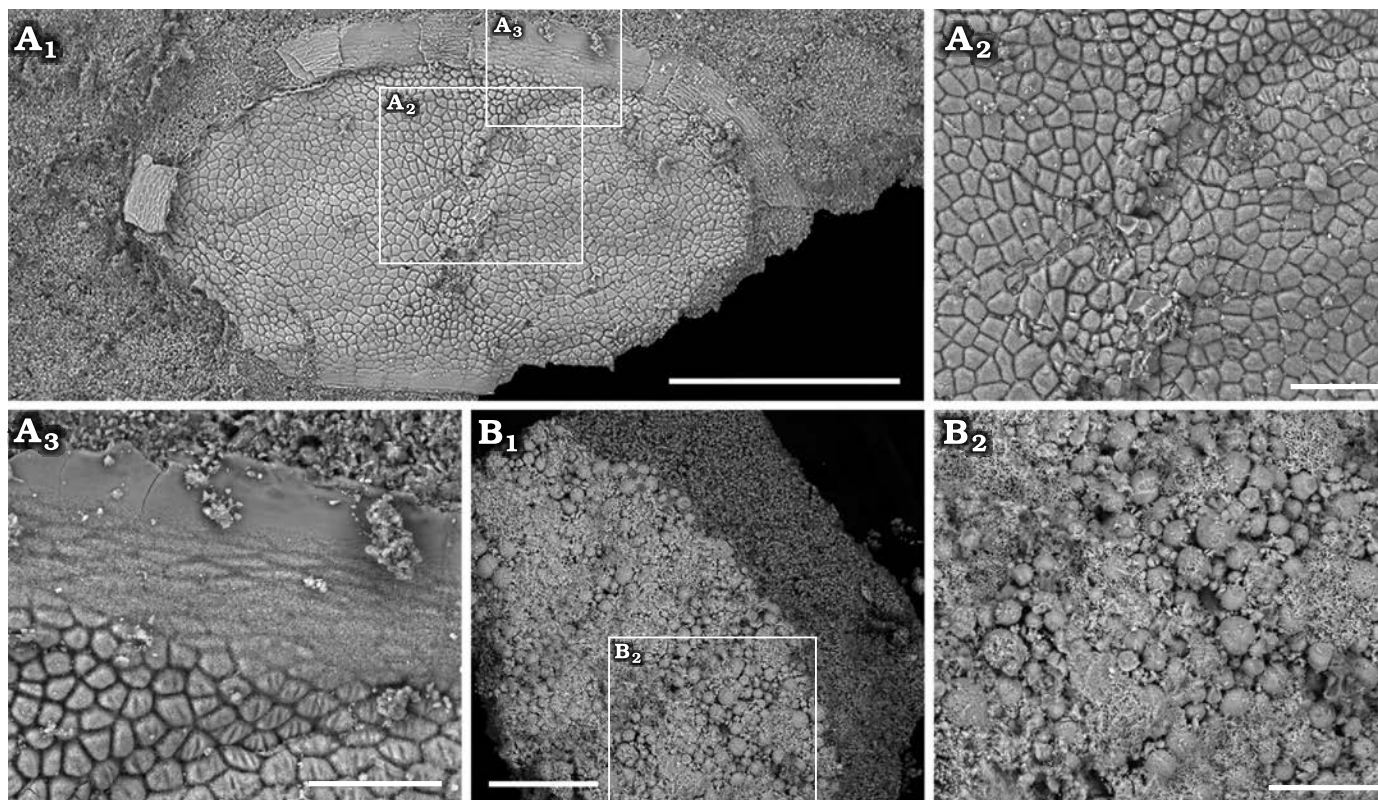


Fig. 5. SEM images of arthropod remains from the lower Miocene Foietta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). **A.** Complete *Daphnia* (*Ctenodaphnia*) sp. (Cladocera) ephippium (A₁) showing an exceptionally well-preserved reticulated microsculpture (FS-43B-4); detail showing the limit between the paired egg chambers (A₂), and the dorsal ridge with variation in microsculpture in the top of the image (A₃). **B.** Chironomid larval gallery (B₁) and its filling material (B₂) resulting from oxidation of the original framboidal pyrite (FS-34A-1). Scale bars: A₁, 0.3 mm; A₂, A₃, B₂, 0.05 mm; B₁, 0.1 mm.

of the Foietta la Sarra-A ostracod shells prevents a proper determination. On the other hand, mass records of cladoceran ephippia occur in the Foietta la Sarra-A slabs (Fig. 4B). Specimens are around 0.8–1 mm long and 0.4–0.5 mm wide with an ovoid or sub-rectangular shape. Although the ephippia are poorly preserved, the margins of the valve (originally highly sclerotised) and the paired egg chambers can be distinguished. The dorsal margin lacks spinules. The ephippia show their typical reticular microsculpture (Fig. 5A) with high value of Ca element (Fig. 6A). The ephippial accumulations do not show uniform orientations in the slabs (see Peñalver et al. 1996). The morphotype of the Foietta la Sarra-A ephippia corresponds to the subgenus *Daphnia* (*Ctenodaphnia*) based on the sub-rectangular shape and the axes of the eggs being sub-parallel to the dorsal margin (Kotov and Taylor 2011), as in two specimens recorded from the San Chils locality (Álvarez-Parra and Peñalver 2019). Miocene cladoceran ephippia have been found in several lacustrine deposits from the Iberian Peninsula, such as Rubielos de Mora (Peñalver et al. 1996), Bellver de

Cerdanya in Lleida (Martín-Closas and Delclòs 2007), and Tresjuncos in Cuenca (Bustillo et al. 2017).

Regarding insects, specimens belonging to the orders Hemiptera, Coleoptera, and Diptera have been found. First, a partly preserved abdomen divided in at least seven sclerites of 1.32 mm in length and 1.46 mm in maximum width has been found (Fig. 4C); its distal area is deformed, and the genitalia are not visible. Its morphology could correspond to a pentatomid hemipteran (Hemiptera: Pentatomidae). This group is represented in the San Chils locality by a *Sciocoris?* sp. specimen (Álvarez-Parra and Peñalver 2019). Moreover, beetles (Coleoptera) are represented among the Foietta la Sarra-A material by a poorly preserved isolated elytron, 0.91 mm in length with rounded apex and ornamentation constituted by ten or 11 rows of hollows (Fig. 4D). Beetles had been previously reported from the Ribesalbes–Alcora Basin (Peñalver et al. 2016; Álvarez-Parra and Peñalver 2019). Lastly, an isolated wing with a rounded apex, six longitudinal veins and a fusiform (discal?) cell likely belongs to a psychodid fly (Diptera: Psychodidae) (Quate and Vockeroth 1981). In

← Fig. 4. Arthropods from the lower Miocene Foietta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). **A.** Undetermined ostracod shell, FS-38A-1. **B.** Mass record of *Daphnia* (*Ctenodaphnia*) sp. (Cladocera) ephippia, FS-21A; each specimen is indicated with an arrowhead. **C.** Photograph (C₁) and drawing (C₂) of a Pentatomidae? (Hemiptera) abdomen, FS-41-1. **D.** Isolated beetle (Coleoptera) elytron, FS-19B-1. **E.** Photograph (E₁) and drawing (E₂) of a Psychodidae? (Diptera) wing, FS-44D-1. **F, G.** Chironomid larval galleries in longitudinal (FS-45-1) and transversal (FS-29) views, respectively. Scale bars: A, B, C, E, F, 1 mm; D, G, 0.5 mm.

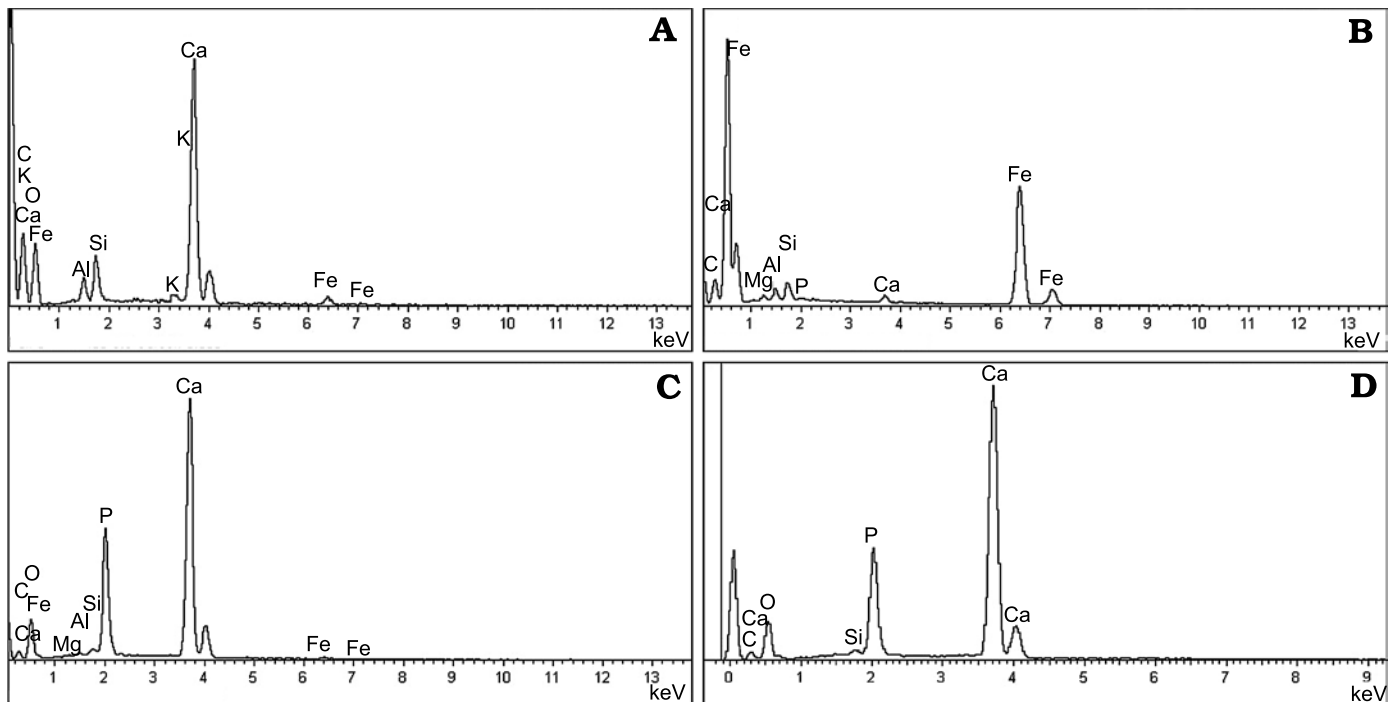


Fig. 6. EDX microanalyses of specimens from the lower Miocene Foieta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). **A.** Ephippial microsculpture of *Daphnia* (*Ctenodaphnia*) sp. (FS-43B-4). **B.** Filling material of a chironomid larval gallery (FS-34A-1). **C.** Morphotype 2 fish scale (FS-21C-9). **D.** Isolated fish bone (FS-21).

any case, this determination should remain tentative as the wing margin is not clearly defined and the veins obscured due to preservation (Fig. 4E). This would be the first report of the family Psychodidae in the Ribesalbes–Alcora Basin. In addition, U-shaped and rather straight galleries of around 0.2–1 mm in diameter and different lengths have been found in the slabs from Foieta la Sarra-A (Fig. 4F); their sections are circular (Fig. 4G). This type of ichnofossils of around 1 mm in diameter is commonly related to bioturbation produced by chironomid (Diptera) larvae and are usually found in lacustrine deposits (Walshe 1951). Similar ichnofossils attributed to chironomid larval galleries have been found in other Cenozoic outcrops from the Iberian Peninsula (e.g., Rodríguez-Aranda and Calvo 1998). The galleries show an orange-brown colour resulted of the oxidation process of the original framboidal pyrite (Fig. 5B), as EDX microanalysis indicates (Fig. 6B).

Fishes.—A poorly preserved and incomplete teleostean specimen, abundant clumps of scales and bones, isolated scales, and other scattered remains comprise the fish record from Foieta la Sarra-A. The incomplete teleostean specimen, preserved as a part and a counterpart, preserved its anterior part. It is comprised by a crushed skull, scales of the abdominal region and the dorsal and pectoral fins (Fig. 7A). The skull is crushed and only the dentary bone and some bones of the opercular apparatus are recognisable, although distinctive features of the fish skull are obscured by dissolution and/or enzymatic attack, which hinder both a detailed anatomic description and an accurate taxonomic assignment. As far as

it can be described, the dorsal fin appears to be triangular; no scutes, basal or fringing fulcra are observed. The dorsal fin base length is about 30 mm and dorsal pterygiophores are badly preserved. There are ca. 15 dorsal fin rays with long bases, in the preserved part of which segmentation and branching are not evident. It is unclear if only the first dorsal fin ray forms the leading margin of the fin. The distal portion of the fin is obscured by sediment. The vertebral column and its associated elements are not apparent. Although, one badly preserved vertebral centrum is observed near the base of the skull, it is not clear whether it belongs to this specimen or not. The squamation is composed by small, thin sub-squarish elasmoid scales of the cycloid type (morphotype 1, see below). Scales are imbricated, most of them completely preserved. Some isolated scales show their anterior field exposed; the posterior field, which is inserted below the epidermis, appears to be large. No lateral line scales have been recognised, nor ctenoid or spinoid scales are observed in the specimen. Circulii are present as concentric crests surrounding the focus area of the scales; the focus seems to be displaced from the central area. Circulii of the posterior and lateral fields are smooth and seems to be regularly spaced. The so-called first circulii are also smooth. The scales are composed by two layers: the superficial layer and the basal bony plate (Fig. 8B). The basal bony plate is the main part of the scale; its structure or size of collagen fibres is not evident in the available material.

The clumps of scale and bones are well-sorted (Fig. 7B, C). Among the bones, some fin rays and ribs have been identified. The scales in the clumps are represented by at least

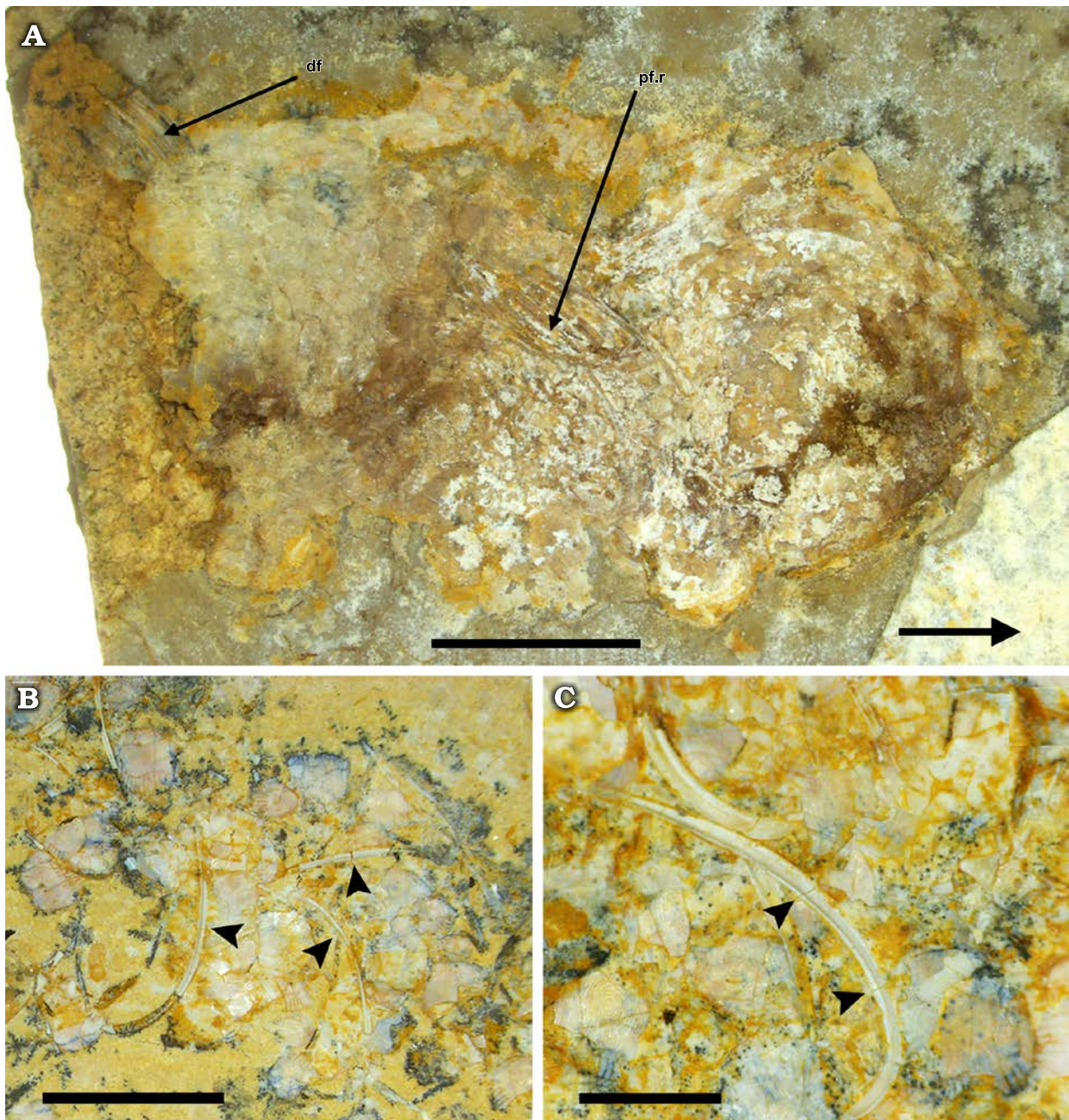


Fig. 7. Fish remains from the lower Miocene Foieta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). A. Poorly preserved teleostean specimen (FS-23A-1); df: dorsal fin, pf.r: pectoral fin rays; the arrow indicates the anterior region of the specimen. B, C. Clump of scales and bones (FS-38A-2); arrowheads indicate diagenetic fractures. Scale bars: A, B, 5 mm; C, 2 mm.

the two morphotypes described below. It is unclear if these two morphotypes of scales belong to the same individual. Moreover, it is also unclear if some of these clumps represent bromalites (i.e., coprolites, cololites, and regurgitalites; see Hunt and Lucas 2012). Some fin rays and ribs (i.e., fractures perpendicular to the major axis of the bone; Fig. 7B, C), as well as most of the scales, exhibit diagenetic fractures.

Fish scales from Foieta la Sarra are represented by at least two morphotypes that are described below. The morphotype 1 is well-represented in the incomplete specimen covered with articulated scales (see above), as well as in the clumps of scales and bones. EDX analysis of fish scales of

morphotype 2 (Fig. 6C) and an isolated bone (Fig. 6D) confirm that the scale and bone are composed namely of Ca, P, and O, indicating the prevalence of apatite.

Morphotype 1: Thin sub squarish elasmoid scales of the cycloid type. These scales measure ca. 1.5 mm long (measured antero-posteriorly) and 1.6 mm wide (measured in the widest portion) (Fig. 8A, B). They have a relatively small focus that is antero-posteriorly elongate. Circuli are well-developed. The anterior field is striated (see Bräger and Moritz 2016: fig. 3B) and bears 13 to 17 radii (= ridges). The so-called first circuli are concave to straight. Lepidonts (= small tooth-like structures anchored to circuli) are not observed.

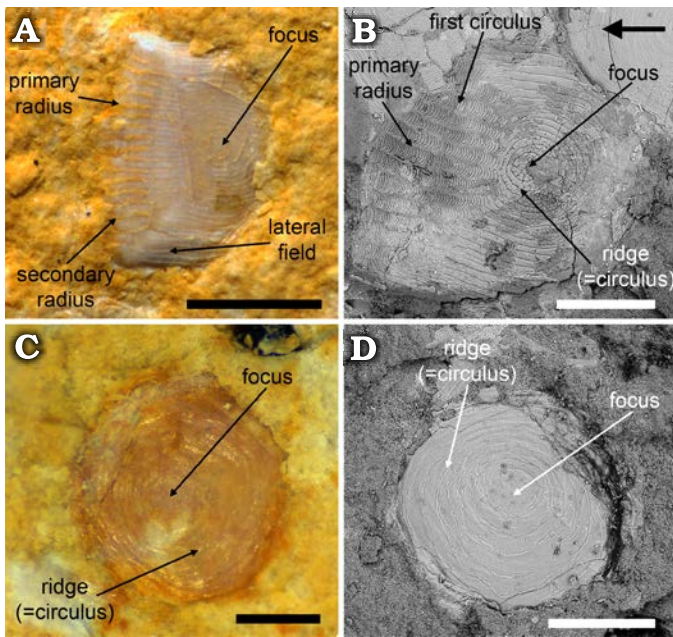


Fig. 8. Fish scale morphotypes from the lower Miocene Foieta la Sarra-A locality (Ribesalbes–Alcora Basin, Castelló Province, Spain). A, B. General view and SEM image of the morphotype 1 (FS-21B-2 and FS-21A-10, respectively); the arrow indicates de anterior field of the scale. C. General view (C₁) and SEM image (C₂) of the morphotype 2 (FS-21C-9). Scale bars: A, 1 mm; B, C, D, 0.5 mm.

The posterior field seems to be convex and smooth. Two generations (principal and secondary) of radius (=ridges) are easily recognised. The posterior field of the scales does not show tubercles, but its absence cannot be fully ascertained due to the preservation of the available material.

This scale morphotype is commonly found in many freshwater fishes (e.g., Cyprinodontiformes and Atheriniformes), and it is most similar to the scales present in some killifish species of the genus *Aphanius* (e.g., Gholami et al. 2013; Teimori et al. 2017).

It is important to highlight that scales might vary in type (ctenoid and cycloid) and morphology (rounded, squarish, etc.) in the same individual relative to the different body sections of an individual. Scales are plastic structures; their general morphology is highly (intra- and interspecifically) variable; also, microstructure seems to show a high degree of intraspecific variation, at least in some cyprinodontiforms such as *Aphanius* (e.g., Gholami et al. 2013) and clupeiforms (e.g., Patterson et al. 2002). Because taxonomy based on isolated scales appears to be problematic (e.g., Bräger et al. 2017), we prefer to leave them unassigned.

Morphotype 2: Elasmoid scale of the cycloid type, thin, rounded or circular in shape, diameter about 1 mm (Fig. 8C, D). The scale has a large focus located in the middle and concentric circuli. Radii and spines or ctenii are lacking. The number of circuli (seven to ten) is almost equal across the entire field. Circuli are almost equally spaced and concentric. This scale resembles those present in the freshwater gadid teleostean *Lota lota* Linnaeus, 1758, which is the only freshwater member of the family Gadidae and is adapted to

cold waters (Lagler 1947; Oates et al. 1993; Daniels 1996). Note that some members of Salmoniformes have similar scale morphology (Patterson et al. 2002).

Discussion

Taphonomy.—The laminated limestone beds from Foieta la Sarra-A are interpreted as lacustrine deposits; the recorded taxa correspond to a typical assemblage of a perennial palaeolake, characterised by the presence of lymnaeid and planorbid molluscs, ostracods, and charophytes (Freyet and Verrecchia 2002). The Foieta la Sarra-A record fulfils the requirements to be considered a Konservat-Lagerstätte based on the preservation of soft-bodied specimens, such as the well-preserved reticulated microsculpture of the *Daphnia* (*Ctenodaphnia*) ephippia (Fig. 5A) and insect remains (Fig. 4C–E), which is explained by the presence of microbial mats in the palaeolake (Wilby et al. 1996). The incomplete fish specimen with articulated bones, fins, and scales in original position (Fig. 7A) could have reached the bottom of the lake articulated in conditions of low water energy (see below). The carcass lacks evidence of flotation, as well as of predation and/or scavenging. Aside from this singular specimen, bacterial decay and flotation appear to be evident in the remaining fish carcasses, as most of the specimens are disarticulated and forming clumps (Fig. 7B, C), perhaps due to scavengers since no clear orientation of the elements is apparent. In any case, at least some of the clumps could be bromalites; thus, their preservation resulted from rapid burial after production. The well-developed diagenetic fractures on bones and scales evidence the fossil diagenesis (Fig. 7B, C). There appears to be no directional scatter of *Daphnia* (*Ctenodaphnia*) sp. ephippia, fish scales, and/or lepidotrichia. Thus, the palaeolake was apparently quiet, lacking currents that could have affected the distribution of fish carcasses. Characite rocks and the slabs with chironomid larval galleries correspond to the in situ record of marginal areas of the lake, while the rest of the specimens were deposited at the lake bottom covered by microbial mats.

The presence of charophytes, mass records of cladoceran ephippia, ostracods, and fish remains in Foieta la Sarra-A indicates that the water chemistry was dissimilar to that of the palaeolake of the Unit B which is apparently lacking these taxa (Peñalver et al. 2016). The laminated slabs from La Rinconada and San Chils resulted from the sedimentation of fine-grained particles on microbial mats at the anoxic bottom from a meromictic lake (Anadón et al. 1989; Peñalver et al. 2016; Álvarez-Parra and Peñalver 2019). The environmental conditions related to the presence of microbial mats in Foieta la Sarra-A could also be different. As noted above, it is unclear if the clumps of fish scales and bones resulted from scavenging or they are bromalites. Therefore, we propose two hypotheses for the presence of microbial mats at the lake bottom: (i) as finding potentially scavenged fish is incompatible with an anoxic lake bottom, the microbial mats were related

to a lake bottom that was at least slightly saline (Bauld 1981; Peñalver and Gaudant 2010); this contradicts the molluscan assemblage found, which indicates freshwater conditions in the palaeolake; (ii) if the fish remains were bromalites, the lake bottom could have been anoxic. The preservation of the Foieta la Sarra-A materials is worse than other Miocene palaeolakes such as Bicorn and La Rinconada (Peñalver and Gaudant 2010; Peñalver et al. 2016), so it is plausible that the microbial mats of Foieta la Sarra-A were thin veils rather than thick mats. A detailed taphonomic study of the Foieta la Sarra-A palaeolake is beyond the aims of this contribution, and future studies will have to address that matter.

Palaeoecological reconstruction of the Foieta la Sarra-A palaeolake and its surroundings.—The botanical and faunal composition of the Foieta la Sarra-A record (Table 1) represents the community that inhabited a lacustrine environment and contiguous areas during the early Miocene.

Charophyte accumulations of *Chara* sp. recorded as characite are related to the presence of charophyte meadows at the lower marginal areas of the lake, which was well-oxygenated and well-illuminated based on the finding of thalli and gyrogonites (Soulié-Märsche et al. 2010). Charophyte stems are well-preserved and somewhat fragmented, suggesting that the environment was relatively quiet, without the action of strong currents nor waves. The charophyte meadows corresponded to one of the primary producers in the palaeolake.

The assemblage of molluscs is composed of 127 poorly preserved specimens of four freshwater gastropod taxa. The 93.3% are pulmonate species (Lymnaeidae indet., *Ferrissia* sp., and *Gyraulus* sp.) which, as a whole, indicate aquatic environments that are shallow, stagnant or with weak currents and abundant aquatic vegetation (Glöer 2002). *Ferrissia* would also indicate the existence of probable reed-belts on the shore while Lymnaeidae and *Gyraulus* point out the possibility of temporary desiccation (Fetcher and Falkner 1993; Glöer 2002). Many extant individuals studied in Europe, determined as *Ferrissia wautieri* (Mirolli, 1960) but according to Vecchioni et al. (2017) corresponding to the allochthonous species of cosmopolitan distribution *Ferrissia californica* (Rowell, 1863), inhabit stagnant to lentic waters, living among the leaves of aquatic vegetation and underneath the leaves of reeds-belts on the upper littoral zone and showing little sensitivity to water quality (Glöer 2002). Some authors also indicate that the same species is thermophilous and lives in eutrophicated waters (Van der Velde 1991). Extant species of Lymnaeidae preferably inhabit shallow, permanently stagnant or weakly flowing waters, with salinities ranging from freshwater to oligohaline in the case of *Stagnicola* and up to mesohaline in *Radix*; lymnaeids typically live on macrophyte vegetation feeding on epiphytes (Adam 1960; Okland 1990; Glöer 2002). Some lymnaeid species withstand periods of desiccation and inhabit temporary water bodies, even those being located on flood plains (Fetcher and Falkner 1993). Extant species of *Gyraulus* live in a variety of conditions, preferably in shallow waters, including

water bodies that are ephemeral, stagnant, or slowly running and abundant in vegetation (Glöer 2002; Welter-Schultes 2012). The fourth taxon (Hydrobiidae? indet.), which represents 6.7% of the specimens studied, suggests permanent waters, at least at the time of deposition. Extant species of Hydrobiidae thrive in permanently flooded freshwater to brackish water environments (Arconada and Ramos 2003).

Lymnaeids feed namely by scraping algae and diatoms from rocks or macrophytes (Pyron and Brown 2015), although they often consume filamentous algae (Kesler et al. 1986). Extant Holarctic species of *Ferrissia* show a preference for periphytic algae, especially diatoms (Burky 1971). Concerning the genus *Gyraulus*, the analysis of stomach content indicates a primarily detritivorous diet (Dudgeon and Yipp 1985), although other studies also show algal consumption (e.g., Hann et al. 2001). The species of Hydrobiidae have a widely diverse diet, including detritus, bacteria, filamentous algae, and diatoms, among others. Studies on the feeding habits of some freshwater hydrobiid species (Radea et al. 2017), as well as species from other related families such as Amnicolidae (Kesler 1981), show a diet suggesting the importance of periphyton diatom grazing. In sum, the combination of the four gastropod taxa found in Foieta la Sarra-A suggest that the palaeolake was shallow, perennial, and quiet (absence of strong currents). Moreover, this molluscan assemblage indicates that the aquatic macrophytes were abundant in the shallow areas of the lake such as the littoral zone, and the shores were also covered with helophytic vegetation. This vegetation was important in maintaining molluscan populations, as constituted the main substrate for the epiphytes on which molluscs fed on, particularly in the case of Lymnaeidae. Moreover, this vegetation would contribute to the formation of detritus which could support the *Gyraulus* sp. populations. These characteristics are compatible with a vegetated shore of the lake whose waters were oligohaline at the most.

Daphnia cladocerans are filter-feeding crustaceans of freshwater habitats (Ebert 2005). The specimens from Foieta la Sarra-A correspond to the subgenus *Daphnia* (*Ctenodaphnia*), which is related to shallow, temporary water bodies (Popova et al. 2016). *Daphnia* (*Ctenodaphnia*) ephippia from Foieta la Sarra-A have been found as mass records in the rock slabs, suggesting environmentally stressful conditions, such as temporary desiccation or lack of food resources, that caused the change in their reproductive cycle from parthenogenetic to ephippial females (Peñalver et al. 1996; Ebert 2005). The finding of these records is interesting, as it appears to contradict the permanent water condition inferred by charophytes, some of the molluscs and fish. Nonetheless, *Daphnia* (*Ctenodaphnia*) individuals could have occupied lake margins that desiccated during dry periods in a lake with a fluctuating extension. Predators of *Daphnia* (*Ctenodaphnia*) could have been other arthropods as well as vertebrates, although their presence did not affect the emergence of massive *Daphnia* populations. Although cladoceran ephippia have not been found in the extensive

fossil record of La Rinconada (Peñalver et al. 2016), and they are represented in San Chils only by two specimens in a slab (Álvarez-Parra and Peñalver 2019), similar ephippial mass occurrences were found in Rubielos de Mora (Peñalver et al. 1996). Moreover, the occurrence of mass records of *Daphnia* (*Ctenodaphnia*) in Foieta la Sarra-A indicates that the water chemistry of this palaeolake was different in comparison to the lacustrine deposits from the Unit B. Furthermore, ephippia from Foieta la Sarra-A do not show a preferential orientation in the slabs, suggesting a low energy environment. Lastly, they are found associated to mass occurrences of ostracods, like the ones from Rubielos de Mora and Bellver de Cerdanya Miocene outcrops (Peñalver et al. 1996).

The absence of dipteran pupal exuviae in Foieta la Sarra-A is noteworthy, as they are abundant in La Rinconada and San Chils, where they indicate stagnant water and lack of predators (Peñalver et al. 2016; Álvarez-Parra and Peñalver 2019). The presence of fish predators matches such absence of dipteran exuviae, although the record of the latter would require a higher grade of exceptional preservation than that observed in Foieta la Sarra-A. Nonetheless, the finding of Chironomidae larval galleries occurring at the lake bottom suggests that populations of this group of flies were present in this lacustrine environment; chironomid larvae are found in ecosystems of different characteristics, although their presence in Foieta la Sarra-A could be related to shallow waters of a lentic environment and were microphagic, i.e., feeding on detritus and small plants or animals (Oliver 1981).

The teleostean fish scale morphotype 1 resembles the scales present in some killifish (Cyprinodontiformes), a large group of secondarily freshwater fishes with a wide geographical distribution. More specifically, the scale morphotype 1 resembles the scales present in species *Aphanius*, including small omnivorous eurythermic and eurhythaline species that inhabit brackish and salt marshes, coastal lagoons, river mouths and freshwater environments (Gaudant 1993; Nelson et al. 2016). Therefore, *Aphanius* species are highly ecologically flexible fishes that tolerate a wide range of temperature and salinities, and so they may occur in a wide range of habitats (Teimori et al. 2014). Despite their omnivory, the diet of some *Aphanius* species has a high degree of seasonal variation and, in general terms, includes several small invertebrates and their eggs, as well as diatoms (Leonardos 2008). Also, some species of *Aphanius* display some differences in habitat preference related to their age (Alcaraz and García-Berthou 2007).

Although the palaeoichthyofauna can be useful to understand the palaeoenvironment, it is important to highlight that “deducing a type of environment from only a single taxon’s autoecology risks falling into a circular reasoning but this analysis is, however valuable if its incorporates a set of signals that point to the same environment” (Cavin 2017: 4). Moreover, due to the high diversity of fishes’ lifestyles and their ability to live in environments with changes in salinity, using them to determine or define types of freshwater environments is challenging.

The palaeovegetation of Foieta la Sarra-A corresponds to a typical riparian assemblage, similar to the La Rinconada and San Chils assemblages related to a lacustrine palaeoenvironment (Barrón and Postigo-Mijarra 2011; Postigo-Mijarra and Barrón 2013; Álvarez-Parra and Peñalver 2019), based on the presence of specimens of Poales, *Dicotylophyllum* sp., and *Salix?* sp. Palaeofires could be recurrent based on the accumulations of fusinite in the Foieta la Sarra-A rock. The scarcity of insects in the fossil record of Foieta la Sarra-A might be explained by the presence of vertebrates such as fishes, which fed on the former when they were washed to the lake. Abundant amphibian, crocodylian (and other reptilian), and mammalian record has been found in the other levels of the Foieta la Sarra section and Campisano ravine localities (Crespo 2017; Crespo et al. 2019c; VDC unpublished data), whereas amphibians and bird (only feathers) remains have appeared in La Rinconada and San Chils (Peñalver et al. 2016; Álvarez-Parra and Peñalver 2019). Therefore, although these vertebrates could have inhabited the riparian environment or that contiguous the palaeolake of Foieta la Sarra-A, there is a gap of such faunal record to date.

Palaeoenvironmental dynamics during the early Miocene in the Ribesalbes–Alcora Basin.—The deposits of the Ribesalbes–Alcora Basin show a dynamic environmental evolution of the water body during the early Miocene. The oil-shale and laminated bituminous dolomicrite of La Rinconada and San Chils localities, belonging to the Unit B, correspond to the record of a meromictic palaeolake (Peñalver et al. 2016). On the contrary, the stratigraphically higher fine grain facies of the Campisano ravine, belonging to the Unit C, represent the distal deposits of an alluvial fan or a muddy floodplain (Crespo et al. 2019c), indicating an environmental change during the early Miocene in the basin. The laminated slabs found in Foieta la Sarra-A hitherto represent the only record of a well-developed palaeolake in the stratigraphy of the Campisano ravine (Crespo et al. 2019c), showing that the palaeoenvironments inferred for the facies of the Unit C are more diverse than previously assumed.

The palaeolake corresponding to the deposits of the Unit B, which outcrops in La Rinconada and San Chils, had a permanent water stratification, with an anoxic lake bottom where microbial mats developed (De las Heras et al. 2003). This water stratification was probably influenced by the inferred subtropical climate that prevailed in the area (Peñalver et al. 2016). Water depth could be several tens of metres in areas away from the shore (De las Heras et al. 2003). The water was basic in character, oligo- or mesosaline and with high sulphate content (De las Heras et al. 2003). The slight salinity could have also contributed to the stratification of the water column (Peñalver et al. 2016).

The Foieta la Sarra-A palaeolake shows taphonomic and palaeoecological similarities to the lacustrine record of the Unit B, such as the fine lamination due to the presence of microbial mats at the lake bottom (which enabled the fossil preservation and is characteristic of Konservat-

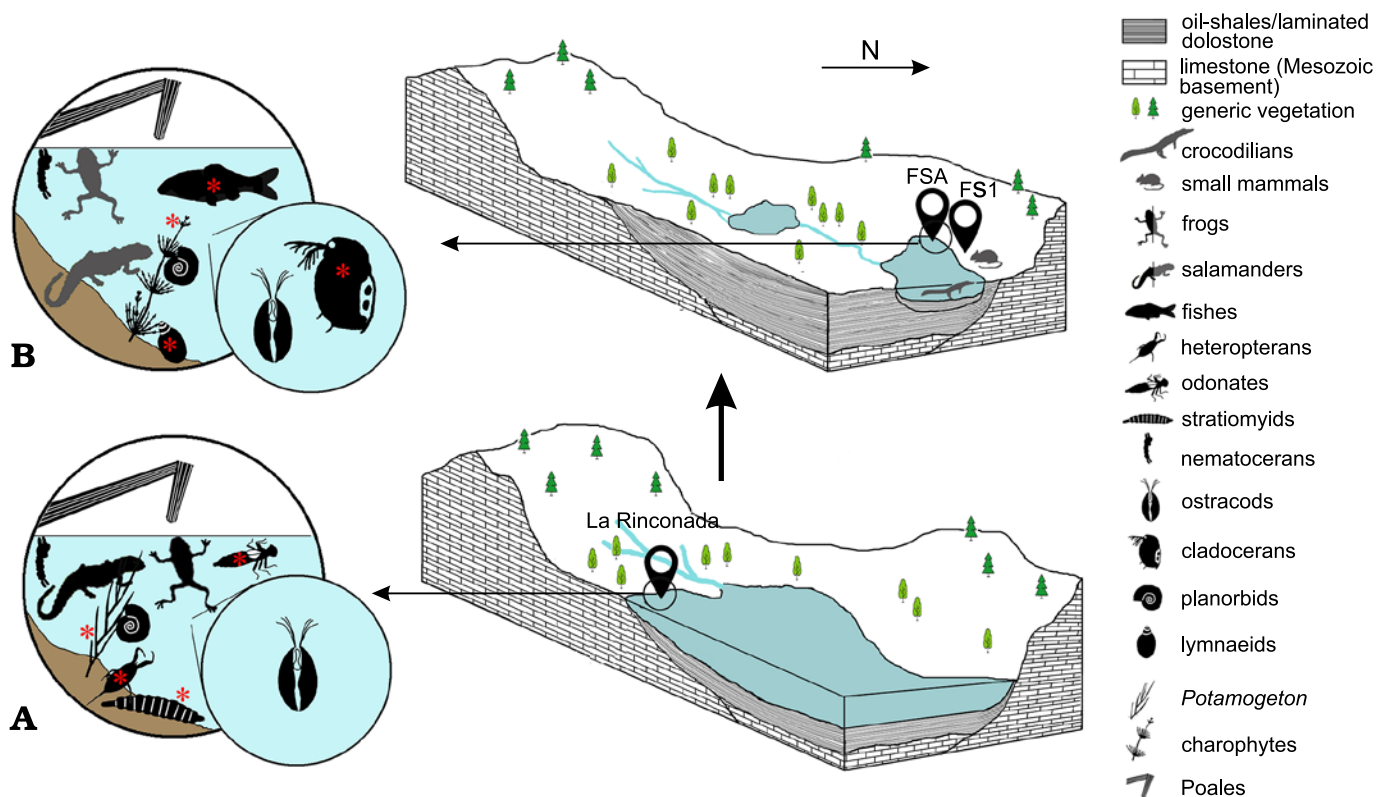


Fig. 9. Synthetic 3D representation of the aquatic environment evolution in the Ribesalbes–Alcora Basin (Castelló Province, Spain) during the early Miocene, mostly based on the La Rinconada (ca. 19 Ma, belonging to the Unit B) and Foieta la Sarra-A (ca. 16.20 Ma, belonging to the Unit C) deposits, which nowadays are separated by around 4 km. Rock deposits have been strongly simplified (e.g., breccia and sandstone are not represented). The significant differences in the aquatic biota are marked with red asterisks. Animals represented in grey correspond to the fossil record of the Foieta la Sarra-1 (FS1) mammalian locality, which are not recorded in the slabs of Foieta la Sarra-A Foieta la Sarra-A (FSA). In La Rinconada (A), heteropterans are represented by the families Nepidae and Notonectidae, odonates by the infraorders Zygoptera and Anisoptera, and nematocerans by the families Chironomidae and Chaoboridae. In Foieta la Sarra-A (B), planorbids are represented by the genera *Ferrissia* and *Gyraulus*, and nematocerans have been inferred based on chironomid larval galleries in the Foieta la Sarra-A slabs. Silhouettes not to scale.

Lagerstätten) and the finding of plant remains and fossil insects (Peñalver et al. 2016). Interestingly, Foieta la Sarra-A shares with San Chils, but not with the widely studied La Rinconada, the presence of *Daphnia* (*Ctenodaphnia*) sp. ephippia, although only represented by two specimens in San Chils (Álvarez-Parra and Peñalver 2019), in contrast to the mass records from Foieta la Sarra-A. La Rinconada has yielded fossil record of amphibians (Peñalver et al. 2016), which have not been found in Foieta la Sarra-A. Despite that, Foieta la Sarra-A has provided charophytes and lymnaeids, not recorded in the Unit B, and the only fish remains of the Ribesalbes–Alcora Basin to date. Considering the overall data from the palaeolakes of the Unit B and Foieta la Sarra-A, the palaeoenvironment and the palaeoecology were clearly different (Fig. 9). Their palaeoenvironments differ due to a change of the water chemistry. The three main mechanisms controlling the water chemistry of lakes are atmospheric precipitation, rock dominance of the basin and evaporation-crystallisation processes (Gibbs 1970), and these three could explain the environmental dynamics of the palaeolake records of the basin. The rock dominance of the basin corresponds to the basement that crops out in the marginal areas of the basin and is eroded by the lotic waters,

which arrive to the lakes formed in the central areas and set the chemistry of their waters (Gibbs 1970). The deposits of the Unit B have been tentatively dated as slightly older than the deposits of the Unit C (Peñalver et al. 2016), but it is unlikely that the rock dominance of the basin could change enough to modify the water chemistry in such a short time. Therefore, the factors that influenced a change of the water chemistry in the palaeolakes of the basin need to be further studied based on sedimentological works and geochemical analyses of the Foieta la Sarra-A rock. An increase of the humidity in the second local biozone of the Campisano ravine was proposed after the preliminary studies of the mammalian palaeoecology and isotopes in the fossiliferous localities (Ríos 2013; Crespo 2017). The faunal assemblage in the second local biozone (after Crespo et al. 2019c) apparently shows more humid preferences than in the first one (Crespo 2017; Crespo et al. 2019c). The Foieta la Sarra-A assemblage and palaeoecology confirm an increase of the humidity in the second local biozone. Indeed, there was a stable water body at least during the beginning of the second biozone, with no lacustrine sediment apparent during the first biozone. A tropical climate has been inferred for the

early Miocene forest of the Campisano ravine surrounding the palaeolake (Crespo et al. 2020b).

The seven sections of the Campisano ravine, corresponding to the Unit C, include facies that represent different palaeoenvironments (Crespo 2017; Crespo et al. 2019c). Most of the deposits are mudstone beds with variegated colours, suggesting that sediments were exposed to subaerial conditions. Sandstone and microconglomerate channels are interpreted as the fluvial contribution to the distal parts of alluvial fans in a low-energy environment. These channels correspond, for instance, to specific high energy episodes related to large floods caused by storms. Thinner tabulate sandstone layers that cover a larger lateral surface, were probably deposited in the floodplains (Crespo 2017; Crespo et al. 2019c). Massive limestone beds were probably deposited in shallow lacustrine and marshy environments subjected to the fluctuation of the water level. Furthermore, several levels of the sections from the Campisano ravine indicate particular environments. These include: (i) dark lenticular beds with black-lignitiferous mudstone probably related to a palustrine environment (in the levels Mas dels Coixos-3 and Mas d'Antolino B-11), (ii) palaeosols associated with longer intervals of subaerial exposure (in the level Mas dels Coixos-4-6), (iii) fine ferruginous and red-coloured levels probably associated to low sedimentation rate episodes and combined with abundant iron supplies (in the level Araia Cantera Sud-2), (iv) evaporitic deposits in the form of gypsum layers formed on the edge of a lake (in the levels Mas d'Antolino B-0A and Mas d'Antolino B-0B), and (v) laminated limestone beds as those from Foietta la Sarra-A (Crespo 2017; Crespo et al. 2019c). Particularly, an increase of the energy in the environment is observed in the Foietta la Sarra section (Fig. 1B), as the laminated limestone beds are at the bottom of the deposit (Foietta la Sarra-A) and sandy mudstone is present at the top (Foietta la Sarra-1). Accordingly, the Campisano ravine deposits reflect the complexity of the palaeoenvironments of a palaeolake and its surroundings. The Foietta la Sarra-A locality in particular shows the environmental evolution of a palaeolake that occurred in the basin in comparison with the record of the Unit B.

Conclusions

Foietta la Sarra-A corresponds to the record of a different palaeolake to that of the Unit B, or a later phase of the same, providing evidence of the palaeoenvironmental dynamics of the Ribesalbes–Alcora Basin during the early Miocene. The Foietta la Sarra-A palaeolake had a different water chemistry in comparison to that of the Unit B. Such factor explains the presence of mass occurrences of ostracods and *Daphnia* (*Ctenodaphnia*) sp. ephippia. The occurrence of characite beds indicates that the palaeolake depth was not significant (i.e., a few metres). Moreover, the water level underwent periodic fluctuations, favouring the mass production of *Daphnia* (*Ctenodaphnia*) sp. ephippia in the most marginal palaeolake areas under stress conditions. The scarcity of

insects could be related to the abundance of fishes as predators, to preservation, and/or to sampling biases.

A new palaeontological work is planned in the near future in order to increase the known fossil assemblage from Foietta la Sarra-A. Furthermore, a sedimentological and geochemical approach would shed further light on the palaeolake environment. The search of new lacustrine limestone levels in the stratigraphy of the Campisano ravine, as well as new palaeontological excavations in La Rinconada and San Chils would provide new data on the complex palaeoecology of the Ribesalbes–Alcora Basin during the early Miocene.

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References

- Adam, W. 1960. *Faune de Belgique. Mollusques. Tome I. Mollusques terrestres et dulcicoles*. 402 pp. Patrimoine de l'Institut royal des Sciences naturelles de Belgique, Brussels.
- Agustí, J., Anadón, P., Ginsburg, L., Mein, P., and Moissenet, E. 1988. Araya et Mira: nouveaux gisements de mammifères dans le Miocène Inférieur-Moyen des Chaînes Ibériques orientales et méditerranéennes. Conséquences stratigraphiques et structurales. *Paleontologia i Evolució* 22: 83–101.
- Alcaraz, C. and García-Berthou, E. 2007. Food of an endangered cyprinodont (*Aphanius iberus*): ontogenetic diet shift and prey electivity. *Environmental Biology of Fishes* 78: 193–207.
- Álvarez-Parra, S. and Peñalver, E. 2019. Palaeontological study of the lacustrine oil-shales of the lower Miocene San Chils locality (Ribe-

- salbes–Alcora Basin, Castellón province, Spain). *Spanish Journal of Palaeontology* 34: 187–203.
- Anadón, P., Cabrera, L., Julià, R., Roca, E., and Rosell, L. 1989. Lacustrine oil-shale basins in Tertiary grabens from NE Spain (Western European rift system). *Palaeogeography, Palaeoclimatology, Palaeoecology* 70: 7–28.
- Arconada, B. and Ramos, M.A. 2003. The Ibero-Balearic region: one of the areas of highest Hydrobiidae (Gastropoda, Prosobranchia, Rissooidea) diversity in Europe. *Graellsia* 59: 91–104.
- Barrón, E. and Diéguez, C. 2001. Estudio macroflorístico del Mioceno Inferior lacustre de la Cuenca de Rubielos de Mora (Teruel, España). *Boletín Geológico y Minero* 112: 13–56.
- Barrón, E. and Postigo-Mijarra, J.M. 2011. Early Miocene fluvial-lacustrine and swamp vegetation of La Rinconada mine (Ribesalbes–Alcora Basin, Eastern Spain). *Review of Palaeobotany and Palynology* 165: 11–26.
- Bauld, J. 1981. Occurrence of benthic microbial mats in saline lakes. *Hydrobiologia* 81: 87–111.
- Bräger, Z. and Moritz, T. 2016. A scale atlas for common Mediterranean teleost fishes. *Vertebrate Zoology* 66: 275–386.
- Bräger, Z., Staszny, Á., Mertzen, M., Moritz, T., and Horváth, G. 2017. Fish scale identification: from individual to species-specific shape variability. *Acta Ichthyologica et Piscatoria* 47: 331–338.
- Burky, A. 1971. Biomass turnover, respiration, and interpopulation variation in the stream limpet *Ferrissia rivularis* (Say). *Ecological Monographs* 41: 235–251.
- Bustillo, M.Á., Díaz-Molina, M., López-García, M.J., Delclòs, X., Peláez-Campomanes, P., Peñalver, E., Rodríguez-Talavera, R., and Sanchiz, B. 2017. Geology and paleontology of Tresjuncos (Cuenca, Spain), a new diatomaceous deposit with Konservat-Lagerstätte characteristics from the European late Miocene. *Journal of Iberian Geology* 43: 395–411.
- Cavin, L. 2017. *Freshwater Fishes: 250 Million Years of Evolutionary History*. 199 pp. ISTE Press, Elsevier, London.
- Cohen, A.S. 2003. *Paleolimnology: The History and Evolution of Lake Systems*. 528 pp. Oxford University Press, New York.
- Costa-Pérez, M., Álvarez-Parra, S., Paredes-Aliaga, M.V., Caballero, Ó., Bueno, E., Vilaplana-Climent, A., and Crespo, V.D. 2019. Valoración patrimonial de los yacimientos del Mioceno inferior del Barranco de Campisano de la Cuenca de Ribesalbes–Alcora (Araia d’Alcora, Castelló, España). *Spanish Journal of Palaeontology* 34: 125–130.
- Crespo, V.D. 2017. *Los mamíferos del Mioceno Inferior de la Cuenca de Ribesalbes–Alcora (Castelló, España)*. 695 pp. Unpublished Ph.D. Thesis, Universitat de València, Valencia.
- Crespo, V.D., Fagoaga, A., Montoya, P., and Ruiz-Sánchez, F.J. 2019a. Oldtimers and newcomers: the shrews and heterosoricids from the Ribesalbes–Alcora Basin (east of Spain). *Palaeontologia Electronica* 22.3.64: 1–22.
- Crespo, V.D., Fagoaga, A., Ruiz-Sánchez, F.J., and Montoya, P. 2021. Diggers, gliders and runners: the squirrels from the Ribesalbes–Alcora Basin (East of Spain). *Bulletin of Geosciences* 96: 83–97.
- Crespo V.D., Gamonal, A., Montoya, P., and Ruiz-Sánchez, F.J. (in press). The eomyids from the Ribesalbes–Alcora Basin (early Miocene, Iberian Peninsula) and their biostratigraphic and paleoecologic implications. *Rivista Italiana di Paleontologia e Stratigrafia*.
- Crespo, V.D., Goin, F.J., Montoya, P., and Ruiz-Sánchez, F.J. 2020a. Early Miocene marsupialiforms, gymnures, and hedgehogs from Ribesalbes–Alcora Basin (Spain). *Journal of Paleontology* 94: 1213–1227.
- Crespo, V.D., Marquina-Blasco, R., Ruiz-Sánchez, F.J., and Montoya, P. 2019b. An unusual insectivore assemblage from the early Miocene of southwestern Europe: the talpids and dimylids from the Ribesalbes–Alcora Basin (Spain). *Comptes Rendus Palevol* 18: 407–416.
- Crespo, V.D., Sevilla, P., Montoya, P., and Ruiz-Sánchez, F.J. 2020b. A relict tropical forest bat assemblage from the early Miocene of the Ribesalbes–Alcora Basin (Castelló, Spain). *Earth and Environmental Science Transactions of the Royal Society of Edinburgh* 111 (4): 247–258.
- Crespo, V.D., Suárez-Hernando, O., Murelaga, X., Ruiz-Sánchez, F.J., and Montoya, P. 2019c. Early Miocene mammal assemblages from the Campisano ravine in the Ribesalbes–Alcora Basin (E Spain). *Journal of Iberian Geology* 45: 181–194.
- Daniels, R. 1996. Guide to the identification of scales of inland fishes of Northeastern North America. *New York State Museum Bulletin* 488: 1–105.
- De las Heras, F.X.C., Anadón, P., and Cabrera, L. 2003. Biomarker record variations in lacustrine coals and oil shales: contribution from Tertiary basins in NE Spain. In: B.L. Valero-Garcés (ed.), *Limnogeology in Spain: a Tribute to Kerry R. Kelts*, 187–228. Consejo Superior de Investigaciones Científicas, Madrid.
- Dudgeon, D. and Yipp, M. 1985. The diets of Hong Kong freshwater gastropods. In: B. Morton and D. Dudgeon (eds.), *Proceedings of Second International Workshop on the Malacofauna of Hong Kong and Southern China*, 491–509. Hong Kong University Press, Hong Kong.
- Ebert, D. 2005. *Ecology, Epidemiology and Evolution of Parasitism in Daphnia*. 98 pp. National Center for Biotechnology Information, Bethesda.
- Fetcher, R. and Falkner, G. 1993. *Moluscos*. 287 pp. Blume, Barcelona.
- Freytet, P. and Verrecchia, E.P. 2002. Lacustrine and palustrine carbonate petrography: an overview. *Journal of Paleolimnology* 27: 221–237.
- Furió, M., Ruiz-Sánchez, F.J., Crespo, V.D., Freudenthal, M., and Montoya, P. 2012. The southernmost Miocene occurrence of the last European herpetotheriid *Amphiperatherium frequens* (Metatheria, Mammalia). *Comptes Rendus Palevol* 11: 371–377.
- García-Paredes, I., Álvarez-Sierra, M.Á., Van den Hoek Ostende, L.W., Hernández-Ballarín, V., Hordijk, K., López-Guerrero, P., Oliver, A., and Peláez-Campomanes, P. 2016. The Aragonian and Vallesian high-resolution micromammal succession from the Calatayud-Montalbán Basin (Aragón, Spain). *Comptes Rendus Palevol* 15: 781–789.
- Gaudant, J. 1993. Un exemple de “régression évolutive” chez des poissons cyprinodontidae du miocène supérieur d’Espagne: *Apanius illunensis* nov. sp. *Geobios* 26: 449–454.
- Gholami, Z., Teimori, A., Esmaili, H.R., Schultz-Mirbach, T., and Reichenbacher, B. 2013. Scale surface microstructure and scale size in the tooth-carp genus *Aphanius* (Teleostei: Cyprinodontiformes) from endorheic basins in Southwest Iran. *Zootaxa* 3619: 467–490.
- Gibbs, R.J. 1970. Mechanisms controlling world water chemistry. *Science* 170: 1088–1090.
- Glöer, P. 2002. *Die Tierwelt Deutschlands, 73. Teil: Die Süßwassergastropoden Nord- und Mitteleuropas. Bestimmungsschlüssel, Lebensweise, Verbreitung*. 327 pp. ConchBooks, Hackenheim.
- Hann, B.J., Mundy, C.J., and Goldsborough, L.G. 2001. Snail-periphyton interactions in a prairie lacustrine wetland. *Hydrobiologia* 457: 167–175.
- Harzhauser, M. and Kowalke, T. 2002. Sarmatian (late middle Miocene) gastropod assemblages of the Central Paratethys. *Facies* 46: 57–82.
- Harzhauser, M., Neubauer, T.A., Gross, M., and Binder, H. 2014. The early middle Miocene mollusc fauna of Lake Rein (Eastern Alps, Austria). *Palaeontographica, Abteilung A: Palaeozoology—Stratigraphy* 302 (1–6): 1–71.
- Hunt, A.P. and Lucas, S.G. 2012. Classification of vertebrate coprolites and related trace fossils. *New Mexico Museum of Natural History and Science* 57: 137–146.
- Kesler, D.H. 1981. Periphyton grazing by *Amnicola limosa*: an enclosure-exclosure experiment. *Journal of Freshwater Ecology* 1: 51–59.
- Kesler, D.H., Jokinen, E.H., and Munns, W.R. 1986. Trophic preferences and feeding morphology of two pulmonate snail species from a small New England pond. U.S.A. *Canadian Journal of Zoology* 64: 2570–2575.
- Kotov, A.A. and Taylor, D.J. 2011. Mesozoic fossils (>145 Mya) suggest the antiquity of the subgenera of *Daphnia* and their coevolution with chaoborid predators. *BMC Evolutionary Biology* 11: 129.
- Kowalke, T. and Reichenbacher, B. 2005. Early Miocene (Ottmangian) Mollusca of the Western Paratethys—ontogenetic strategies and palaeoenvironments. *Geobios* 38: 609–635.
- Lagler, K.F. 1947. Lepidological Studies 1. Scale Characters of the Families of Great Lakes Fishes. *Transactions of the American Microscopical Society* 66: 149–171.
- Leonardos, I. 2008. The feeding ecology of *Aphanius fasciatus* (Valenciennes, 1821) in the lagoonal system of Messolongi (western Greece). *Scientia Marina* 72: 393–401.

- Mandic, O., Hajek-Tadesse, V., Bakrač, K., Reichenbacher, B., Grizelj, A., and Miknić, M. 2019. Multiproxy reconstruction of the middle Miocene Požega palaeolake in the Southern Pannonian Basin (NE Croatia) prior to the Badenian transgression of the Central Paratethys Sea. *Palaeogeography, Palaeoclimatology, Palaeoecology* 516: 203–219.
- Martin-Closas, C. and Delclòs, X. 2007. The Miocene paleolake of La Cerdanya (Eastern Pyrenees). *Geo-Guias* 3: 181–205.
- McNamara, M.E., Orr, P.J., Alcalá, L., Anadón, P., and Peñalver, E. 2012. What controls the taphonomy of exceptionally preserved taxa—environment or biology? A case study using frogs from the Miocene Libros Konservat-Lagerstätte (Teruel, Spain). *Palaios* 27: 63–77.
- Nelson, J.S., Grande, T., and Wilson, M.V.H. 2016. *Fishes of the World*. Fifth edition. 707 pp. Wiley, New Jersey.
- Neubauer, T.A., Mandic, O., and Harzhauser, M. 2011. Middle Miocene freshwater mollusks from Lake Sinj (Dinaride Lake System, SE Croatia; Langhian). *Archiv für Molluskenkunde: International Journal of Malacology* 140: 201–237.
- Neubauer, T.A., Mandic, O., and Harzhauser, M. 2013. The Middle Miocene freshwater mollusk fauna of Lake Gacko (SE Bosnia and Herzegovina): taxonomic revision and paleoenvironmental analysis. *Fossil Record* 16: 77–96.
- Neubauer, T.A., Harzhauser, M., Georgopoulou, E., Kroh, A., and Mandic, O. 2015a. Tectonics, climate, and the rise and demise of continental aquatic species richness hotspots. *Proceedings of the National Academy of Sciences* 112: 11478–11483.
- Neubauer, T.A., Harzhauser, M., Kroh, A., Georgopoulou, E., and Mandic, O. 2015b. A gastropod-based biogeographic scheme for the European Neogene freshwater systems. *Earth-Science Reviews* 143: 98–116.
- Neubauer, T.A., Mandic, O., and Harzhauser, M. 2015c. The freshwater mollusk fauna of the middle Miocene Lake Drniš (Dinaride Lake System, Croatia): a taxonomic and systematic revision. *Austrian Journal of Earth Sciences* 108: 15–67.
- Oates, D.W., Krings, L.M., and Ditz, K.L. 1993. *Field Manual for the Identification of Selected North American Freshwater Fish by Fillets and Scales*. 181 pp. University of Nebraska, Lincoln.
- Okland, J. 1990. *Lakes and Snails. Environment and Gastropoda in 1500 Norwegian Lakes*. 515 pp. Universal Book Service/Dr. W. Backhuys, Oegstgeest.
- Oliver, D.R. 1981. Chironomidae. In: J.F. McAlpine, B.V. Peterson, G.E. Shewell, H.J. Teskey, J.R. Vockeroth, and D.M. Wood (eds.), *Manual of Nearctic Diptera, Vol. 1, 27 (29)*, 423–458. Research Branch Agriculture Canada, Ottawa.
- Patterson, R.T., Wright, C., Chang, A.S., Taylor, L.A., Lyons, P.D., Dallimore, A., and Kumar, A. 2002. Atlas of common squamatomological (fish scale) material in coastal British Columbia and an assessment of the utility of various scale types in paleofisheries reconstruction. *Palaeontologia Electronica* 4: 1–88.
- Peñalver, E. 2002. *Los insectos dípteros del Mioceno del Este de la Península Ibérica; Rubielos de Mora, Ribesalbes y Bicorp. Tafonomía y sistemática*. 548 pp. PhD Thesis. Universitat de València, Valencia.
- Peñalver, E. and Gaudant, J. 2010. Limnic food web and salinity of the upper Miocene Bicorn palaeolake (eastern Spain). *Palaeogeography, Palaeoclimatology, Palaeoecology* 297: 683–696.
- Peñalver, E., Barrón, E., Postigo Mijarra, J.M., García Vives, J.A., and Saura Vilar, M. 2016. *El paleolago de Ribesalbes. Un ecosistema de hace 19 millones de años*. 201 pp. Servicio de Publicaciones, Diputació de Castelló and Instituto Geológico y Minero de España, Ministerio de Economía y Competitividad, Castellón de la Plana.
- Peñalver, E., Martínez-Delclòs, X., and De Renzi, M. 1996. Registro de pulgas de agua [Cladocera: Daphniidae: *Daphnia* (*Ctenodaphnia*)] en el Mioceno de Rubielos de Mora (Teruel, España). *Comunicaciones de la II Reunión de Tafonomía y Fossilización* 1: 311–317.
- Popova, E.V., Petrusek, A., Kořínek, V., Mergeay, J., Bekker, E.I., Karabanov, D.P., Galimov, Y.R., Neretina, T.V., Taylor, D.J., and Kotov, A.A. 2016. Revision of the Old World *Daphnia* (*Ctenodaphnia*) *similis* group (Cladocera: Daphniidae). *Zootaxa* 4161: 1–40.
- Postigo-Mijarra, J.M. and Barrón, E. 2013. Zonal plant communities of the Ribesalbes–Alcora Basin (La Rinconada mine, eastern Spain) during the early Miocene. *Botanical Journal of the Linnean Society* 173: 153–174.
- Pyron, M. and Brown, K.M. 2015. Introduction to Mollusca and the Class Gastropoda. In: J.H. Thorp and D.C. Rogers (eds.), *Thorp and Covich's Freshwater Invertebrates, Ecology and General Biology, Fourth Edition, 18*, 383–421. Academic Press, London.
- Quate, L.W. and Vockeroth, J.R. 1981. Psychodidae. In: J.F. McAlpine, B.V. Peterson, G.E. Shewell, H.J. Teskey, J.R. Vockeroth, and D.M. Wood (eds.), *Manual of Nearctic Diptera, Vol. 1, 27 (17)*, 293–300. Research Branch Agriculture Canada, Ottawa.
- Radea, C., Louvrou, I., Konstantinos Bakolitsas, K., and Economou-Amilli, A. 2017. Local endemic and threatened freshwater hydrobiids of Western Greece: elucidation of anatomy and new records. *Folia Malacologica* 25: 3–13.
- Ríos, M. 2013. *Estudio multi-isotópico de la paleoecología y la paleoclimatología de la Cuenca de Ribesalbes–Alcora (Castellón, España) durante el Óptimo Climático del Mioceno*. 76 pp. Unpublished MSc Thesis, Universitat de València, Valencia.
- Rodríguez-Aranda, J.P. and Calvo, J.P. 1998. Trace fossils and rhizoliths as a tool for sedimentological and palaeoenvironmental analysis of ancient continental evaporite successions. *Palaeogeography, Palaeoclimatology, Palaeoecology* 140: 383–399.
- Schultze, H.-P. 2015. Scales, enamel, cosmine, ganoine, and early osteichthyans. *Comptes Rendus Palevol* 15: 83–102.
- Soulié-Marsche, I., Bieda, S., Lafond, R., Maley, J., Baitoudji, M., Vincent, P.M., and Faure, H. 2010. Charophytes as bio-indicators for lake level high stand at “Trou au Natron”, Tibesti, Chad, during the Late Pleistocene. *Global Planetary Change* 72: 334–340.
- Talbot, M.R. and Allen, P.A. 1996. Lakes. In: H.G. Reading (ed.), *Sedimentary Environments: Processes, Facies and Stratigraphy*, 5–36. Blackwell Publishing, Oxford.
- Teimori, A., Esmaeli, H.R., Erpenbeck, D., and Reichenbacher, B. 2014. A new and unique species of the genus *Aphanius* Nardo, 1827 (Teleostei: Cyprinodontidae) from Southern Iran: a case of regressive evolution. *Zoologischer Anzeiger* 253: 327–337.
- Teimori, A., Motamedi, M., and Manizadeh, N. 2017. Microstructural characterization of the body key scale morphology in six Iranian endemic *Aphanius* species (Cyprinodontidae): their taxonomic and evolutionary significance. *Journal of Ichthyology* 57: 533–546.
- Van der Meulen, A.J., Paredes, I.G., Sierra, M.A., van den Hoek Ostende, L.W., and Hordijk, K. 2012. Updated Aragonian biostratigraphy: small mammal distribution and its implications for the Miocene European Chronology. *Geologica Acta* 10: 159–179.
- Van der Velde, G. 1991. Population dynamics and population structure of *Ferrissia wautieri* (Mirolli, 1960) (Gastropoda, Ancyliidae) in a pond near Nijmegen (The Netherlands). *Hydrobiological Bulletin* 24: 141–144.
- Vasilyan, D. 2020. Fish, amphibian and reptilian assemblage from the middle Miocene locality Gračanica—Bugojno palaeolake, Bosnia and Herzegovina. *Palaeobiodiversity and Palaeoenvironments* 100: 437–455.
- Vecchioni, L., Marrone, F., Aculeo, M., and Arizza, V. 2017. Are there autochthonous *Ferrissia* (Mollusca: Planorbidae) in the Palaearctic? Molecular evidence of a widespread North American invasion of the Old World. *The European Zoological Journal* 84: 411–419.
- Vilanova y Piera, J. 1859. Memoria geognóstico-agrícola sobre la provincia de Castellón. *Memorias de la Real Academia de Ciencias de Madrid* 4: 577–803.
- Walshe, B.M. 1951. The feeding habits of certain chironomid larvae (subfamily Tendipedinae). *Proceedings of the Zoological Society of London* 121: 63–79.
- Wautier, J. 1975. Présence d'Espèces du genre *Ferrissia* Walker, 1903 (Gastropoda, Basommatophora) dans le Néogène du bassin rhodanien (France). *Geobios* 8: 423–433.
- Welter-Schultes, F. 2012. *European Non-marine Molluscs, A Guide For Species Identification*. 760 pp. Planet Poster Editions, Göttingen.
- Wilby, P.R., Briggs, D.E., Bernier, P., and Gaillard, C. 1996. Role of microbial mats in the fossilization of soft tissues. *Geology* 24: 787–790.