Exceptional preservation of Cretaceous biota (plants and invertebrates) in flint-bearing alterites from Moragne (western France)

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ABSTRACT

Cretaceous fossil inclusions were recently discovered in silica-rich nodules from the diachronic alteritic deposits of the Moragne hill, Charente-Maritime, western France. Nodules contain marine organisms such as sponge spicules, putative algae, echinoids (*Periaster undulatus* Agassiz and Desor) and rudists (*Sphaerulites foliaceus* Lamarck; hippuritids and radiolitids). These inclusions co-occur with fragmented pinnae of ferns as well as diverse structures of conifers including leafy axes (*Geinitzia* sp.; *Glenrosa carentonensis* Moreau, Néraudeau, Tafforeau and Dépré) and isolated ovulate scales or subtending bracts of cones. The palaeontological assemblage shows that the alterites from Moragne are condensed deposits resulting from the alteration of the Cenomanian to Turonian-Senonian regional series. Conifer leafy axes display diverse kinds of preservation: permineralization in silica, empty external casts of the gross morphology, and casts filled by ferruginous siltstone. Propagation phase-contrast X-ray synchrotron microtomography revealed exceptional preservation of some leafy axes up to the cell level. We support a multi-phase silicification including at least: (1) a precocious phase occurring during the sedimentation of the Upper Cretaceous deposits or during the earliest diagenesis, and that could be linked with the abundance of sponge spicules; then (2) a late phase occurring during Eocene-Oligocene period that corresponds to a hard climate event with strong and long paedogenetic alteration associated with an important phase of silicification (nodulation). The alterites from Moragne are an uncommon example of long-term time-averaged deposits caused by compaction and selective dissolution of matrix in the context of a late continental weathering.
INTRODUCTION

To determine the age and origins of flint-bearing alterites is commonly problematic because they can result from the erosion and condensation of diverse geological units, and are sometimes diachronic (Dewolf, 1970; Daniou, 1982, 1984). Locally, such superficial deposits contain silica-rich nodules that yield abundant fossil inclusions whose stratigraphic origins and preservation are precious tools to discuss the genesis of alterites. In France, whereas Cretaceous marine invertebrates (brachiopods, bivalves, bryozoans, echinoids, sponges) are common inside flint from alterites (e.g., Néraudeau 2004, 2011; Caux, 2015), terrestrial floras remain extremely rare. Over the last decades, alterites containing fossiliferous flints with Upper Cretaceous plant macroremains were discovered from few localities in western France: the Font-Benon quarries in Charente-Maritime (Moreau et al., 2014a); Claix and Torsac in Charente (Néraudeau, 2014; Moreau et al., 2016); and three areas around Châtellerault in Vienne (Moreau et al., 2018). By contrast with most of Cretaceous plant beds from western France which preserved foliar remains as impressions or compressions with or without cuticle (e.g., Lecointre and Carpentier, 1938; Alvarez-Ramis et al., 1981; Berthelin and Pons, 1999; Néraudeau et al., 2005, 2012; Saint-Martin et al., 2013; Valentin et al., 2014), flints may contain exquisite siliceous preservations of the plant macroremains up to the cell levels.

Recently, flint nodules containing Cretaceous terrestrial plant inclusions and marine invertebrates were discovered in alterites from the village of Moragne, in Charente-Maritime. In order to discuss age and origin of the alterites, we describe the palaeontological assemblage of nodules and its preservation. X-ray synchrotron microtomography is used to determine the degree of preservation of plant macroremains and to characterize their histology.

GEOGRAPHICAL AND GEOLOGICAL SETTING

The study area is located in the northern part of the Aquitaine Basin at Moragne, 14 km east of Rochefort (Charente-Maritime department, southwestern France). The Moragne area consists of a small hill of lower Cenomanian sandstone and clay, which are overlain by Palaeogene alteritic deposits (Bourgueil et al., 1972; Figure 1). Regionally, Cretaceous deposits lie unconformably on the Late Jurassic limestone deposits that constitute the outcrops around the Moragne hill (Figure 1). The fossiliferous flints studied here were all collected in the Eocene-Oligocene residual formation that constitutes the top of the hill and is mainly composed of gravel, sand, silt, and clay. At Moragne, the formation yielding fossiliferous flints are not well developed being limited to a small area of less than 800 m in diameter and 50 cm to some metres in thickness. Alterites yield abundant yellow, orange, red to brown flints.

MATERIALS AND METHODS

Plant inclusions from the fossiliferous flints of Moragne were first observed by one of us (ED) in 2013. Thousands of flints were examined and hundreds of fossiliferous flints were collected on surfaces of agricultural fields between 2013 and 2018. Only some of them bear plant inclusions. Plant remains consist of fragmented pinnae of ferns, leafy axes and isolated scales of cones of conifers, and wood (Figures 2-5). Wood fragments are found isolated in alteritic deposits whereas leafy axes are present on the broken surfaces or inside flints. Size of wood fragments varies from several centimetres
Propagation phase-contrast X-ray synchrotron microtomography (PPC-SRμCT) was used on the best preserved plant specimens and was efficient to characterize their histology. The flint nodules were characterized at beamline ID19 of the European Synchrotron Radiation Facility (ESRF, Grenoble France) using PPC-SRμCT. The setup consisted of: filtered white beam with an effective energy of 125 keV (wigglter W150b gap 35 mm, 6 mm copper filter); sample-detector propagation distance of 100 cm; an indirect detector comprising a 200 μm LuAG scintillator, 3x magnification from a set of Hasselblad lenses and a PCO edge 4.2 detector, producing data with an isotropic voxel size of 2.26 μm. The acquisition consisted of 6000 projections (0.01s exposure time) over 360º, with an offset on the centre of rotation to increase the lateral field of view. The reconstruction was performed with PyHST2 (Mirone et al., 2014), using single distance phase retrieval (Paganin et al., 2002) and an unsharp mask to limit the induced blurring (Sanchez et al., 2012). Data were finally converted into a 16 bit tiff stack of slices (0.001% exclusion of the 32 bits 3D histogram), corrected for residual ring artefacts (Lyckegaard et al., 2011) and cropped. The three-dimensional (3D) reconstructions of specimens were achieved with the software VG Studio Max 2.2 (Volume Graphics, Heidelberg, Germany). The virtual dissection and segmentation were made using 3D region growing and manual selections.

SYSTEMATIC PALAEOBOTANY

Order FILICALES Bower, 1899
Filicales indet. "Fern Type 1"
Figure 2

Material. 1 specimen, IGR-PAL-2782.
Description. The specimen displays three distal fragments of pinnae. The largest pinna fragment is 7 mm long and 5 mm wide. The pinna consists of a rachis bearing sub opposite, scythe-shaped, up to 3.5 long and 2 mm wide pinnules (Figure 2). They are attached by a broad base to the rachis and display an acute apex and an entire margin. The rachis is less than 1 mm wide.
Remarks. Ferns correspond to the less abundant remains in the silicified flora from Moragne. Because of a limited preservation, the veins cannot be distinguished.

Order CONIFERALES Gorozhankin, 1904
Genus GEINITZIA Endl., 1847
Geinitzia sp.
Figure 3

Description. The longest specimen is 70 mm. Branches arise at up to 45°. Shoots are up to 18 mm in diameter, straight to slightly curve. Leaves are helically arranged. They are elongated, up to 21 mm long and 2.5 mm wide (Figure 3). Leaves show a long free part that forms an acute angle of up to 70° with main axis of the shoot. The leaf margin is entire. The apex of leaves is acute and slightly curved inward. In cross section, leaves are quadrangular in shape.

Remarks. Leafy axes of Geinitzia are the most abundant plant remains from the silicified flora of Moragne. Elongated, lanceolate, helically arranged leaves forming an acute angle with the main axis of the shoots are known among several Cretaceous conifers such as Cunninghamites Presl in Sternberg, 1838, Elatocladus T.Halle, 1913, Geinitzia and Pagiophyllum Heer emend. T.M.Harris, 1979. The genus Geinitzia differs from Elatocladus by the absence of contraction at the leaf base (Kvaček, 1999). Leaves of Geinitzia differ from Cunninghamites in having a triangular to quadrangular form in cross section, not needles flattened (Kvaček, 1999; Herman and Kvaček, 2010).

Glenrosa carentonensis Moreau, Néraudeau, Tafforeau and Dépré, 2015

Figure 4

Material. 1 specimen IGR-PAL-2804.

Description. Leafy axis is unbranched, 10.0 mm long and 4.5 mm wide (Figure 4.1-3). Leaves are helically arranged, short, up to 3 mm long and up to 1.5 mm wide. They are triangular in abaxial and adaxial views, rhomboidal in transversal section, claw-shaped in sagittal section (Figure 4.4-6). Their margin is entire. Apex of leaves is obtuse (Figure 4.1-6). Abaxial and adaxial surface of leaves bear putative deep stomatal crypts (Figure 4.7). Crypts are generally as wide as deep, or slightly deeper than wide. Putative stomatal crypts are up to 80–110 μm in diameter and 60–140 μm deep (Figure 4.7). Cuticle is only partially preserved been often torn and up to 13 μm thick. Adaxial epidermis is 12–40 μm thick (Figure 4.8). Ordinary epidermal cells are square, rectangular, or polygonal, being 10–55 μm of wide. Palisade parenchyma consists of one layer (rarely two) of elongated cells (Figure 4.4). Palisade parenchyma occurs on both sides but is mainly developed on the adaxial side. It develops along the entire leaf length. Cells are 35–110 μm long and 30–60 μm wide. Anticlinal walls are curve to straight. Spongy parenchyma is well-developed both on abaxial and adaxial sides (Figure 4.4-6). Spongy parenchyma is thinner toward to the leaf apex and absent on this latter. Spongy parenchyma consists of large and rounded cells. Cells vary in size, being 15–130 μm in diameter. The anticlinal walls are curve. Transfusion tissue occurs on the abaxial side and is up to 200 μm thick (Figure 4.4-6). Transfusion tracheids are slightly elongate and ovoid to polygonal. Transfusion tracheids are 25–100 μm long and 10–45 μm wide. A single vascular bundle ends before the leaf apex (Figure 4.4-6). Vascular bundle forms an angle of up to 45° with the stem. Vascular bundle is up to 1.5 mm long, up to 120 μm wide (Figure 4.4-6). Vascular bundle is basally embedded in transfusion tissue.

Remarks. Due to the poor preservation of some parts of the specimen and technical limits of the tomographic protocol, papillae from crypts (see Moreau et al., 2015), stomata, hypodermis, and details of vascular cylinder are not accessible. The genus Glenrosa is represented by seven species that are Barremian to Turon-Coniacian in age (e.g., Watson and Fisher, 1984; Srinivasan, 1992; Zhou et al., 2000; Moreau et al., 2015; Néraudeau et al., 2016). Glenrosa carentonensis highly differs from other species in developing a phyllotaxy 8/21, claw-shaped leaves, a thicker cuticle, and a higher number of papillae and stomata per crypt.

Coniferales indet. “Scale/bract Type 1”

Figure 5

Material. 1 specimen IGR-PAL-2795c

Description. The scale/bract is longer than wide (i.e., 6.5 mm long and 5.0 mm wide), convex,
shows a broad truncate base, an acuminate apex and an entire margin (Figure 5.1-2). The apex displays a free tip (2 mm long and up to 1 mm wide) that bends upward. The visible adaxial surface displays a central, proximal, 3.0 mm long, and 0.5 mm wide, groove. The groove ends before the apical free tip.

Remarks. Due to a limited preservation, it is not clear that the specimen is an ovulate cone scale rather than its subtending bract. Some isolated ovulate cone scales preserved as impressions and compressions were previously reported from the lower Cenomanian of Charente-Maritime (e.g., Néraudeau et al., 2009). They were tentatively...
FIGURE 4. Flint from Moragne containing an Upper Cretaceous conifer leafy axis Glenrosa carentonensis which displays a shoot with helically arranged and short leaves, IGR-PAL-2804. 1-2, Photographs of the broke-open nodule and detail of the leafy axis. 3-9, X-ray synchrotron microtomography virtual histological sections showing preservation of all inner tissues; longitudinal section of the leafy axis (3); sagittal section of a leaf (4); transversal section of a leaf (5); tangential section of a leaf (6); uppermost adaxial surface of a leaf showing a putative stomatal crypt in transversal section (7); uppermost adaxial surface with cuticle, epidermis and hypodermis in transversal section (8); detail of vascular bundle and transfusion tracheids in sagittal section (9). Scale bars: 1, 1 cm; 2, 0.5 mm; 3, 2 mm; 4-6, 0.5 mm; 7-9, 0.2 mm. Abbreviations: Cu., cuticle; Ep., Epidermis; P.pa, palisade parenchyma; P.sp, spongy parenchyma; St.c, putative stomatal crypts; Tr.t, transfusion tracheids; Va., vascular bundle. Voxel size: 3-9, 2.26 μm.
ascribed to *Alvinia bohemica* (Velenovský) Kvaček, 2000. However, the shape of the cone scale from Moragne strongly differs from these specimens. We notice that the gross morphology and the range size of the specimen from Moragne are close to those of the some bract scales of the Cretaceous Cheirolepidiaceae (e.g., Del Fueyo et al., 2008).

**OTHER ORGANISMS**

The most abundant marine invertebrate remains correspond to sponge spicules, with dozens to thousands of specimens preserved in a same flint nodule. Fragmented or complete rudists are the second most frequent silicified invertebrates. Fragments of the Cenomanian rudist *Sphaerulites foliaceus* Lamarck, 1819, are particularly abundant in the alterites but never found in the flints (Figure 6.1). In addition, four sub-complete rudists have been found in the lateritic deposits. The two best ones can be identified as large sub-cylindrical *Hippurites* sp. or *Radiolites* sp. (Figure 6.2-4), species that characterize deposits younger than Cenomanian in Charentes (Macé-Bordy, 2007; Néraudeau et al., 2013b). Five colonial corals close to *Eosiderastrea* sp. have been collected in the alterites, outside the flints (Figure 7.1). A single solitary coral, close to *Aulosmilia* sp. has been found (Figure 7.2). Three specimens of the large Cenomanian spatangoid echinoid *Periaster undulatus* Agassiz and Desor, 1847 have been collected, two in reddish flint nodules, the last one in the alterites (Figure 7.3-5).

Finally, the silicified wood from Moragne is sometimes perforated by large and long galleries likely made by the xylophagous bivalve *Teredo* sp. These galleries have sometimes a blue opalescent colour that contrasts with the reddish oxidized colour of the fossil wood (Figure 6.5). The flints from Moragne yielded a single specimen of putative alga that superficially resemble *Solenopora* Dybowski, 1877 (Figure 8).

**DISCUSSION**

**Age of the Fossil Assemblage**

*Geinitzia* and *Glenrosa* were reported in many Albian to Turonian plant beds from western France (e.g., Néraudeau et al., 2005, 2009, 2016; Girard et al., 2013; Néraudeau, 2014; Moreau et al., 2015, 2017). *Geinitzia* was previously reported from the uppermost Albian of the Font-Benon and Les Renardières quarries (Charente-Maritime; Néraudeau et al., 2005; Moreau et al., 2017), the Cenomanian of the Puy-Puy and Le Brouillard quarries (Charente-Maritime and Maine-et-Loire respectively; Néraudeau et al., 2005, 2013a) and from the Turonian of Simeyrols (Dordogne; Saint-Martin et al., 2013). In all these localities, specimens are preserved as compressions in marls with lignite. Only the Cenomanian flints of Archingeay-Les Nouillers (Moreau et al., 2014a), and those from the Turono-Coniacian of Claix previously yielded permineralized/petrified leafy axes of *Geinitzia*. In western France, cuticles of the conifer *Glenrosa* were previously reported from many Albian to Turono-Coniacian deposits from Aude, Charente, Charente-Maritime and Dordogne (see Moreau et al., 2017 and reference therein). After the silicified deposits from the Font-Benon quarries (Charente-Maritime), the flints from Moragne constitute the second occurrence worldwide yielding both *Geinitzia* and *Glenrosa* preserved as siliceous permineralization (Moreau et al., 2014a, 2014b, 2015).

The echinoid *Periaster undulatus* characterizes the upper part of the first marine lithological unit (B1 sensu Néraudeau et al., 1997; Néraudeau and Moreau, 1989) of the lower Cenomanian from northern Aquitaine (Néraudeau, 2012). The rudist *Sphaerulites foliaceus* is known in the lower and mid Cenomanian deposits from Charentes (Néraudeau et al., 2013b; Miramand et al., 2017). However, the hippuritid and radiolitid rudists indicate a younger age, late Turonian to Senonian.

Thus, although flints from the Moragne hill were previously and tentatively dated to Cenomanian (e.g., Laporte et al., 2000), this study demonstrates that the alterites consist of condensed deposits, which result from the alteration of Creta-
FIGURE 6. Upper Cretaceous silicified marine bivalves from the flints of Moragne. 1, Fragment of the Cenomanian rudist *Sphaerulites foliaceus*, IGR-PAL-2805. 2, Lateral view of radiolitid rudist, IGR-PAL-2806. 3, Lateral view of an hippocritid rudist, IGR-PAL-2807. 4, Top view of an hippocritid rudist showing the two alveoles, with quartz geodes, that correspond to the two teeth of the opercular valve, IGR-PAL-2808. 5, Piece of silicified wood with two galleries realized by the xylophagous bivalvia *Terebo* sp., IGR-PAL-2809. Scale bars equal 1 cm.
aceous formations at least lower Cenomanian to Turonian-Senonian in age (Figure 9). At Moragne, several dozens of metres of the original Upper Cretaceous series finally correspond to only less than one to some metres of alteritic deposits bearing flints (Figure 9).

**Origin of the Alterites**

As well as in Moragne, other localities from Charente and Charente-Maritime show silicified deposits at the top of topography. It consists of diachronic silicification affecting all deposits independently to their age and lithology, from the Upper Jurassic to the lower-middle Eocene (e.g., Néraudeau, 2011; Daniou, 1978). The Eocene-Oligocene period is characterized by a hard climate episode with strong and long paedogenetic alteration associated to an important phase of silicification. Regionally, the phenomenon was tentatively explained by a kaolinization of surface deposits (see Moreau et al., 2014a, 2014b). Exposed rocks were dissolved and decalcified under the influence of soil leaching by meteoric waters during warm and wet climates (Gallois, 2009). As explained by Callen (1983) the dissolution and the re-deposition of silica commonly result from a climate change from wet to arid. Regionally, the top of alterite deposits yields flints containing upper Eocene-Oli-
Although a main silicification event occurred obviously during the late Palaeogene, it is not sufficient to explain the exquisite preservation of tissues and cells in 3D inside Late Cretaceous fossil plants. Such preservation is only possible when the mineralization of inner and intracellular spaces occurs precociously. The abundance of sponge spicules in the matrix of plant-bearing flints as well as on the surface of plant remains (see Figure 10) might have been a potential source of silica. Thus, the silicified fossil assemblage from the flints of Moragne supports a multi-phase silicification including: (1) at least a precocious phase occurring during the sedimentation of the Upper Cretaceous deposits or during the earliest diagenesis; (2) a late phase occurring during Eocene-Oligocene interval and explaining the nodulation leading to secondary flints. Such multi-phase silicification processes have been previously identified in several localities yielding plant-bearing flints from western France (Moreau, 2014b, 2017). The various kinds of plant preservation (empty casts, replicas by siltstone, permineralizations) among plant macro-remains show that silicification occurred heterogeneously, probably depending on the concentration of silica-rich sponge spicules in the diverse sediments. Empty casts and replicas of leafy axes are not compressed but preserved in 3D. It confirms that plant remains, which were dissolved during the
main phase of weathering, were mineralized (at least partially) during the earliest stages of the diagenesis.

Deposits showing mixing of non-contemporaneous fossils in a single bed are called "time-averaged deposits" (Walker and Bambach, 1971). As demonstrated by many authors (e.g., Fursich and Aberhan, 1990; Kidwell and Holland, 2002; Bush et al., 2002), such deposits are commonly observed in marine environments. However, they are less documented in context of continental weathering. Commonly, the main factors causing time-averaged deposits are biological (e.g., biogenic reworking and bioturbation) and sedimentological (currents and low sediment input produce erosion, winnowing, reworking, bypassing of sediment and transport). Here, the alterites from Moragne is an uncommon example of long-term time-averaged deposits caused by compaction and selective dissolution of matrix during late weathering period mainly controlled by climatic events.

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