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Approach to the knowledge of preservation of pleistocenice bone: The case of a Gomphotherium cranium from the site of Tepeticpac, Tlaxcala, Mexico

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ABSTRACT:

An almost complete cranium of a gomphotherium found in Tepeticpac, Tlaxcala, was analyzed with X-ray diffraction (XRD), Petrography and Scanning Electron Microscopy with Energy Dispersive Spectroscopy (SEM-EDS) to establish the transformations

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of the bone during its burial. The analyses assessed that the bone mineral (bioapatite) had suffered modifications in mineral composition, "crystallinity index", cell parameters and CO₂ content. However, the paleohistological structure of the bone was not significantly affected, although evidence of microbial attack was found in the bone surfaces. The filling minerals of bone macro and micro porosity were also analyzed; its main filling minerals are calcite (micrite and sparite), clays, and minerals of volcanic origin.

KEYWORDS: Bone, gomphotere, mineralization, X-ray diffraction, petrography.

RESUMEN:

Un cráneo casi completo de gonfoterio encontrado en Tepeticpac, Tlaxcala, se estudió mediante difracción de rayos X, petrografía y microscopía electrónica de barrido con espectroscopía de dispersión de rayos X para establecer las transformaciones que sufrió el hueso durante su enterramiento. Estos análisis permitieron determinar que el mineral del hueso (bioapatita) experimentó modificaciones en cuanto a su composición mineral, índice de cristalinidad, parámetros de celda y contenidos de CO₂. No obstante, la estructura paleohistológica del hueso no se transformó de manera significativa, aunque se encontraron evidencias de ataque biológico en las áreas superficiales de las muestras de hueso. Los minerales de relleno de la macro y microporosidad del hueso también se caracterizaron y se comprobó que los minerales más abundantes son calcita (micrita y esparita), arcillas de tipo expansible y minerales de origen volcánico.

PALABRAS CLAVE: Palehistología, hueso, gonfoterio, mineralización, difracción de rayos X, petrografía..

INTRODUCTION

An almost complete skull of a proboscide was found in Tlaxcala in 2014. The remains were identified by Dra. Maria Teresa Alberdi as a gomphothere (*Cuvieronius* sp.) based on the morphology of the dentition and the spiral-shaped band of tusks enamel.

The cortical bone, better preserved than the trabecular bone, showed heterogeneity in color, density and porosity. Several analyses have been carried out to understand the bone conservation and the postmortem bone mineral transformations (diagenesis and possible fossilization processes) with the aim to propose better preservation methodologies for the cranium. This remain is now located in the Coordinacion Nacional de Conservacion del Patrimonio Cultural for its restoration and then it will be at display at the Museo Regional de Tlaxcala.

Pleistocene megafauna remains are quite frequent in Mexico (Ferrusquia-Villafranca *et al.*, 2010). The primary interest in paleontological remains has been taxonomic (*e.g.* Garcia-Zepeda and Garduno-Monroy, 2006; Robles-Camacho *et al.*, 2010), however, fossils were often recollected without following a systematic procedure, in which stratigraphic context and sedimentology were overlooked. The relationship between initial peopling and megafauna in the Mexican territory was also documented (Cruz-y-Cruz *et al.*, 2015; Arroyo- Cabrales *et al.*, 2006; Vinas-Vallverdu *et al.*, 2015). More recently, the isotopic signatures and stratigraphy of these paleontological remains have been used as paleoenvironmental records (Pate *et al.*, 1989; Wings, 2004; Robles-Camacho *et al.*, 2010; Perez-Crespo *et al.*, 2012; Cruz-y- Cruz *et al.*, 2016; Perez-Crespo *et al.*, 2016).

The *in vivo* bone structure is composed of an organic and inorganic fraction forming a complex material that provides flexibility and strength to the skeleton. The organic part is formed mainly by collagen. The inorganic, or mineralized, fraction is composed of microcrystalline bioapatite, always very similar in structure to hydroxylapatite but differing in composition by the substitution of carbonate for PO₄³⁻ (biopatite in fresh bone can contain 7% wt. CO₃²⁻), the presence of minor elements into lattice, and water incorporation (Wopenka and Pasteris, 2005).

During early diagenetic processes, the soft tissues are removed, collagen is degraded and mineralization, including chemical changes like fluor (F) enrichment and structural changes (*e.g.* cell parameters) transform bioapatite into a more thermodynamically stable phase. Apatite can incorporate a big number of chemical elements and the stoichiometric and perfect hexagonal structure is only possible for the structure of fluorapatite (Sponheimer and Lee-Thorp, 1999; Hedges, 2002; Jans, 2008).

Apatite is a mineral group with complex solid solutions. In order to simplify, the International Mineralogical Association (IMA) accepted minerals considering three end members: hydroxylapatite, fluorapatite and chlorapatite (Chang *et al.*, 1996; Back, 2014). The different size of monovalent anions (F^- , Cl^- , OH^-) determines a variation in cell parameters. The informal names francolite and dahllite have been applied for apatites enriched in CO_2 and fluorine, and apatites enriched only in CO_2 , respectively (Deer *et al.*, 2013).

The study of fossil diagenetic processes from neoformed and precipitated minerals in the vascular cavities of the bone and in the fractures allows to establish the physical and chemical conditions that existed in the lithosphere during lithification, permineralization, and other fossilization processes (Casal *et al.*, 2017). Thus, it is important to evaluate the conservation state, post mortem alteration and approximately estimate the impact of the transformations in its geochemical and isotopic signatures.

DESCRIPTION OF THE STUDY AREA

On May 2014, workers of the Tepeticpac Archaeological Project (TAP) of the Instituto Nacional de Antropología e Historia (INAH), accidentally uncovered around 10 centimeters of the distal section of a gomphothere tusk (*Gomphotheriidae* family). Afterwards, controlled excavations were undertaken using an archaeological exploration methodology in order to provide adequate stratigraphic information of local deposits. An area of four square meters was excavated using a horizontal grid made of four 1 x 1 m quadrants. Vertical depth was registered using a level bank placed over the deposits in the UTM coordinates: 0579830, 2138859, at an elevation of 2,298 meters above sea level (Figure 1). If we project the finding coordinates in the interactive geological map published by Ferrari *et al.* (2018) we can verify that the location where the gomphothere cranium was found correspond to the unit described as: Volcano-Sedimentary and Lacustrine Deposits (Late Miocene to Pleistocene).

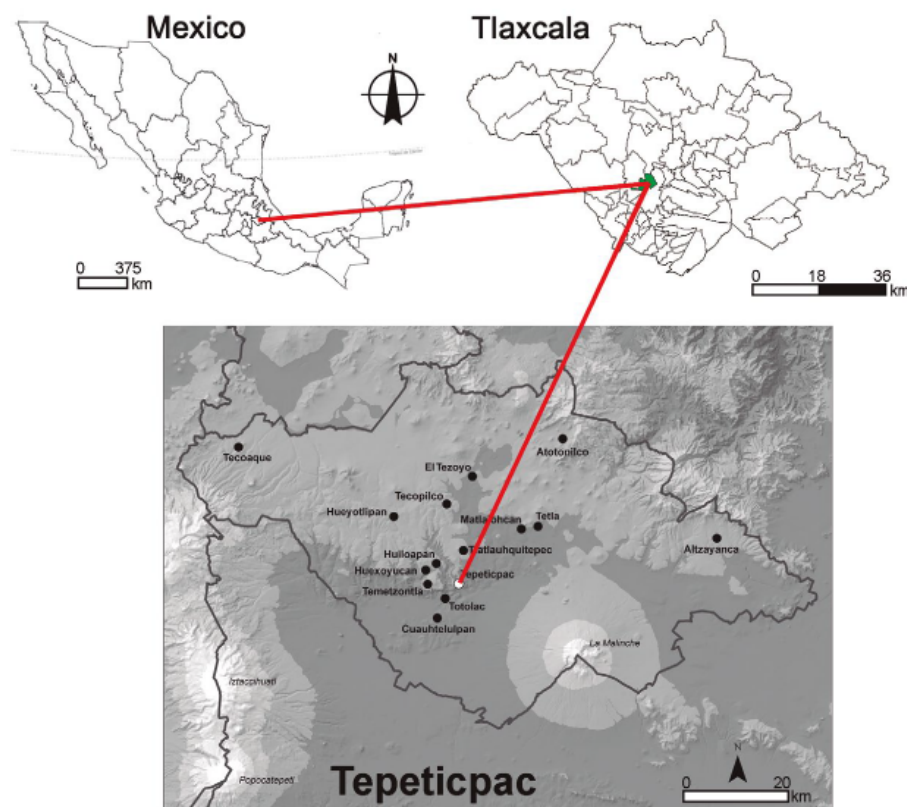


FIGURE 1
Location Site.

The discovery took place on an alluvial deposit exposed by a crosscutting road on the hillslopes of Tenextepetl hill, which is part of a mountain chain system that elevates around 300 meters above the Puebla-Tlaxcala valley and that separates the bottom of the valley to the south from the elevated Tlaxcala Block to the north.

The uppermost stratum was formed of 20 to 30 centimeters of loosened tepetate rock that had a steep north to south inclination, which matches the present day-slope surface. It contained no paleontological remains. Below the upper stratum lays Layer A, a colluvial stratum consisting of blocks of medium size blocks (10 to 20 cm) of calcareous rocks and fragments of tepetate rock (fragipan-indurated tephra material). The deposit also had a north to south slope and a thickness of around 1 meter; it was strongly compacted, although some sand lenticular deposits were observed on the north and west profiles.

Below Layer A, on the north portion of the excavation, we detected Layer B, a deposit formed by the superimposition of several thin layers of fine to coarse rounded sands, some of them forming diagonal bands. These seemed to be part of alluvial or lacustrine deposits. In some sections, the sands were interspersed with thin layers of carbonates. The sand layers had grayish tones (10YR 6/1) and covered more than half of the excavation.

Layer C was the bottommost sediment. It is an alluvial stratum made up of medium to coarse rounded sands, probably corresponding to old lacustrine sequence, and very similar in composition to Layer B, but without carbonates. This layer extended throughout the excavation and had a very gentle inclination from north to south, discordant with the steep present day slope. It included dispersed concretions and small intrusions of Layer A materials, mainly calcareous rocks and medium size (10 to 20 cm) blocks of tepetate rock, particularly near the east profile of the exploration.

Correlating the studied section with the complete Pliocene- Quaternary paleosol-sedimentary sequence of Tlaxcala block (Sedov *et al.*, 2009) we relate the B and C layers with the uppermost part of the uplifted

Pliocene lacustrine sedimentary unit. Colluvial Layer A incorporates materials of the calcareous Pliocene lake deposits as well as fragments of Pleistocene tepetate. Six to eight layers of tepetate (indurated tephra) interlayered with paleosols formed during the middle and late Pleistocene constitute a semi-continuous mantle over the Tlaxcala Block (Heine and Schonhals, 1973; Sedov *et al.*, 2009). As far as colluvial Layer A includes redeposited materials derived from both Pliocene (lacustrine) and Pleistocene (tephra-paleosol) units, we suppose that its development post-dates these units. We associate this colluviation with the phase of high geomorphic activity and intensification of surface erosion/sedimentation processes which occurred in the Central Mexican Highlands in the Terminal Pleistocene (late Glacial) (Heine, 1984; Sedov *et al.* 2009). This chronostratigraphic interpretation supposes a major time gap (*hiatus*) between formation of the Layer A and layers B and C.

Excavation of the skeletal remains was extremely difficult due to the compactness of the stratum, and by intrusive remnants of Layer A. At around 50 to 60 centimeters of excavation, we uncovered the top portion of the gomphothere cranium, which was comprised of the parietal, frontal and zygomatic bones, along with its intact right tusk and a partially looted left one. The cranium had its frontal bones upwards, and had an east-west orientation with its tusks pointing towards the west. Part of the front portion of the skull had collapsed due to the weight exerted by the upper layers. The tusks, maxillae and teeth were well-preserved. The rest of the cranium bones, especially the frontal bone, were fragile and fragmented due to compression because medium sized rocks that laid on top heavily smashed them. Some sections of the bones were absent such as portions of the zygomatic. Also, parts of the cranium bones were partially covered with a fine layer of clayish material of Layers A and B. It appears that sediments of Layer A incorporated and sealed the upper half of the fossil, while the lower section intruded into the underlying B and C layers (Figure 2).

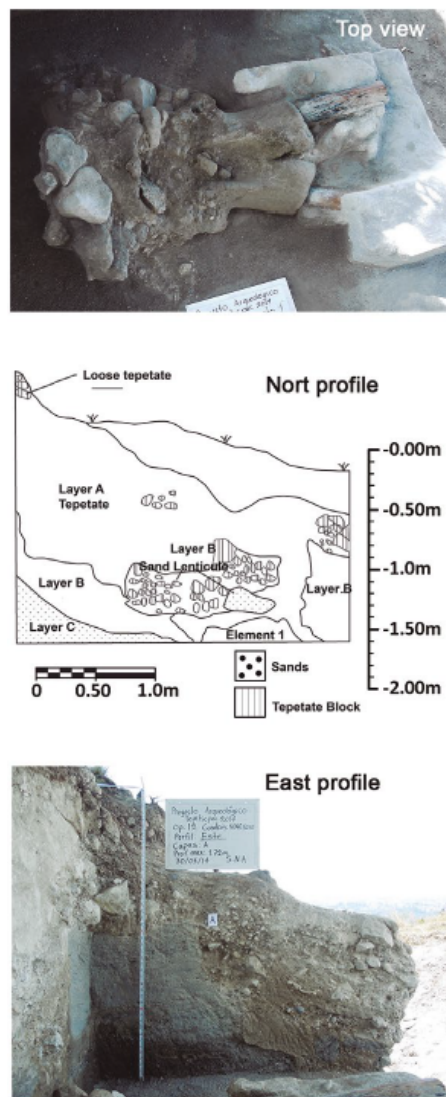


FIGURE 2
North and east profiles with a top view of the cranium.

MATERIALS

Three samples of bone were used for thin sections: a) Cortical bone and trabecular bone with crystalline minerals in the walls of the trabecula partially filled; b) Trabecular bone totally filled with sediments; c) Fractured trabecular bone cemented with the sediments of the filling (Figure 3). A total of six thin sections, two per sample were obtained, however, only one uncovered thin section from sample 2 was analyzed in SEM-EDS.

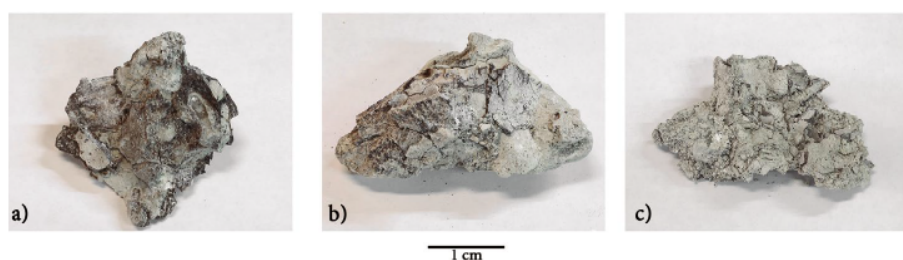


FIGURE 3

a) Cortical bone and trabecular bone with crystalline minerals in the walls of the trabecula partially filled. b) Trabecular bone totally filled with sediments. c) Fractured trabecular bone cemented with the sediments of the filling. We obtained two thin sections of each sample.

Samples of detached cortical bone, that was better conserved than the trabecular, were classified in three categories according to their anatomical position, color and porosity: 1- frontal bone with a white yellowish color in the anterior face, the internal side of the cortical bone has a brownish color, they both have low porosity and diagenetic alterations such as pitting and delamination; the pittings and surfaces losses were filled with a white material; 2- temporal bone with a brownish yellow color with more porosity and irregularities in the superficial area of the bone, the pores are filled with white and opaque mineral and with a translucent mineral. The internal face is black; 3- alveolus, the bone has a darker color and has more pores and other cavities which are filled with a white mineral. The internal face is black (Figure 4).

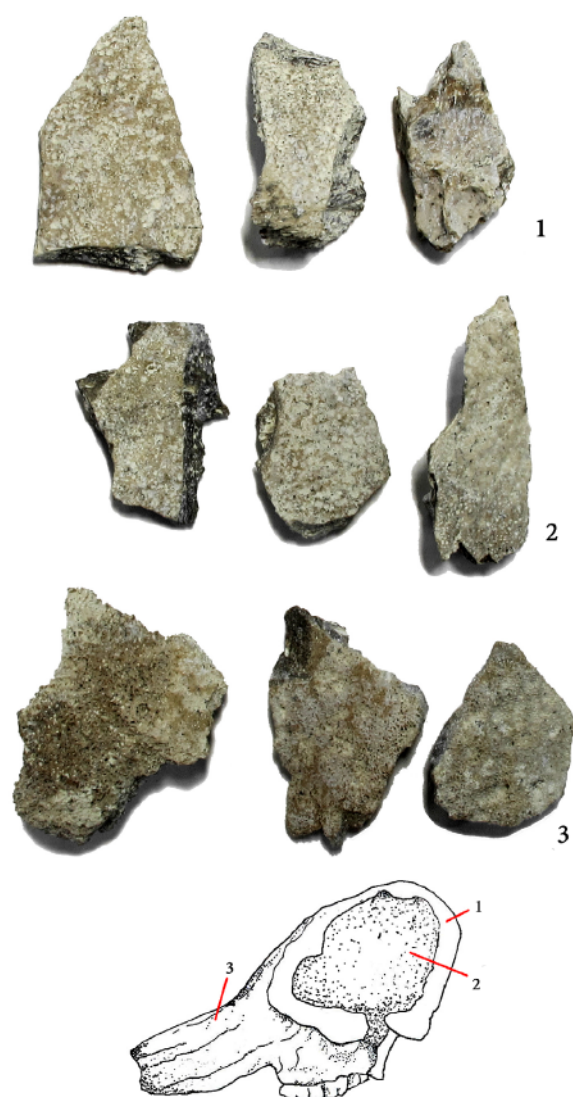


FIGURE 4

Cortical bone used for DRX, the sample sizes are no more than 3 cm long. 1) Less porous samples, 2) samples with medium porosity, 3) samples with the most porosity.

A detailed mineralogical study, using Powder X-ray Diffraction (PXRD) and Rietveld refinement, allow to characterized the differences between the tree categories to understand the permineralization of the bones and to identify the presence of diagenetic processes. For comparison purposes (Figure 5), we also use a pure inorganic well crystalized apatite (Cerro de Mercado, Durango fluorapatite), an actual bone of low crystallinity (cow bone composed by hydroxilapatite). We also measure with PDRX three samples associated with the sediments that covers the skeleton.

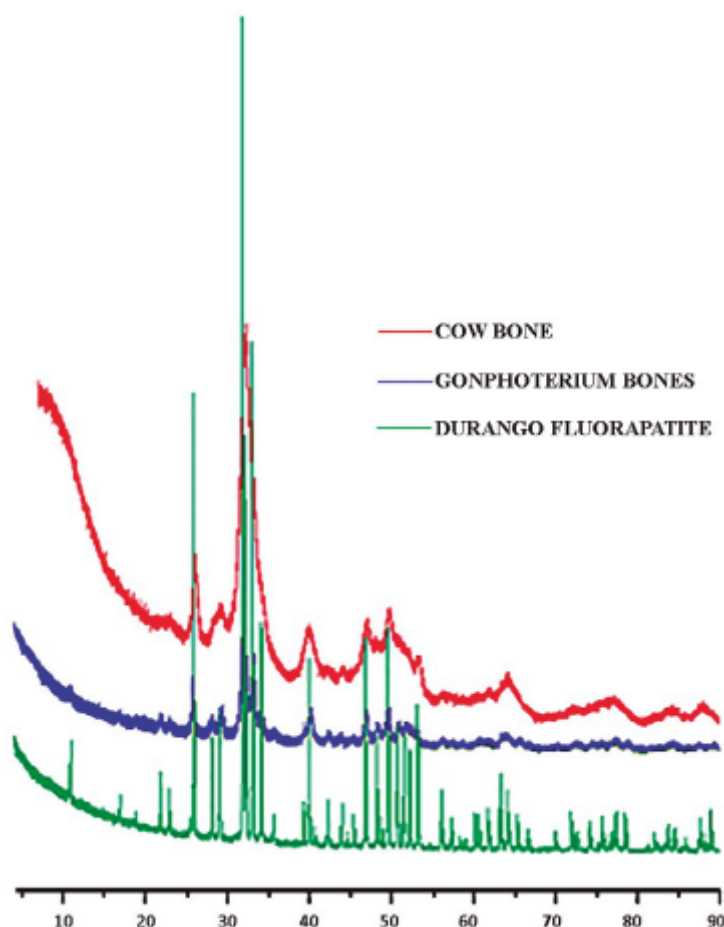


FIGURE 5

X-ray patterns of the Gomphoterium analyzed samples (blue) with intermediate crystallinity and the pure inorganic (Cerro de Mercado fluorapatite) and pure organic (cow bone hydroxylapatite) end members.

METHODS

Petrography

Thin sections allowed us to analyze the micromorphology of cortical and trabecular bone as well as the distribution and composition of subsequent layers of sediments. Histological analyses has been extensively used in archaeological and paleontological researches (Chinsamy and Raath, 1992; Hubert *et al.*, 1996; Chinsamy, 1997; Jans, 2008; Lamm, 2007; Guido *et al.*, 2012; Casal *et al.*, 2017; Tomassini *et al.*, 2015, Tomassini *et al.*, 2014; Pfretzschner, 2004).

Cortical and trabecular bone with their sediment fillings were encapsulated, hardened and immobilized with a polyester resin applied with vacuum. Then the obtained blocks were cut with a diamond cutter until reaching almost 30 μm . The section is adhered to a sample carrier and then thin sections were polished by hand. An Olympus Bx51 petrographic microscope with digital camera attached and with several types of light (plain polarized light (ppl), cross polarized light (xpl) and reflected light (rfl)) was used. Analyses were performed using the Image Pro Puls 5.1 software.

SEM/EDS

Scanning Electron Microscopy along with an Energy Dispersive X-ray Spectroscopy (SEM - EDS) were used to relate the micromorphology of the bones and the chemical variability of the samples including some diagenetic features, such as tunneling, in the bone fragments surface (Hubert *et al.*, 1996; Reiche *et al.*, 2003; Guido *et al.*, 2012).

Thin section of an uncovered sample 2 along with a non-treated sample in a transversal section of cortical bone were considered for this analysis. A JEOL JSM6060LV with an INCAEnergy 250 EDSLKIE250 were used, under the following parameters: 20 kv, vacuum, back scattered signal, and diverse magnifications.

X-ray diffraction

X-ray diffraction (XRD) was used to determine the mineral composition of the bone, and to calculate several crystallographic parameters that help us to understand the bone mineral transformations, such as “crystallinity index”, crystallite size, parameters a and c of unit cells and CO₂ percentage (Person *et al.*, 1995; Bocherens *et al.*, 2008; Piga *et al.*, 2009; Peters *et al.*, 2000; Reiche *et al.*, 2003; Keenan *et al.*, 2015).

Sample Preparation

The bone samples were ground with an agate pestle and mortar to <75 µm and mounted in aluminum backside holders for X-ray powder diffraction analysis.

Clay Sample was gently disaggregated to avoid artificial grain size reduction of rock components, then was broken into small chips (2 mm) using a porcelain crusher and subsequently dispersed in deionized water. Clay size fraction (<2 µm) was separated in distilled water according to Stoke’s law using the most unaggressive method (Moore and Reynolds, 1997).

From the <2 µm fraction, air-dried oriented preparation was obtained by pipetting some drops of the suspensions onto a glass slide, which was then dried at 30 °C for a few hours (Moore and Reynolds, 1997). Ethylenglycol solvation of the slide was achieved by exposing them to ethylenglycol vapor at 70 °C for a minimum of 12 hours.

Measurement

Measurements were carried out on a PANalytical Empyrean diffractometer equipped with a PIXcel3D detector and nickel filter, operating with an accelerating voltage of 45kV and a filament current of 40mA, using Cu Kα radiation. All bone samples were measured over a 2θ angle range of 4–90 °, with a step size of 0.02° (2theta) and 40 s of scan step time.

Clay sample was examined by XRD in the air-dried form, saturated with ethylene glycol (EG) and after heating (550 °C). The preparations were measured over a 2θ angle range of 4–70 ° (air-dried) and 4–30 . (glycolated and heated).

Phase identification and characterization

Rietveld refinement (Rietveld, 1967) was applied (using Highscore 4.5 PANalytical software) to calculate cell parameters and “XRD Crystallinity index” (Person *et al.*, 1995). Crystallite size and CO₂ content (O’Brien

et al., 1990) were calculated to determine if the textural features observed in the petrographic study can be confirmed in a quantitative form using PXRD.

In XRD the degree of crystallinity is measured considering the separation of [202], [300], [211] and [112] reflections, located between 30° and 35° for a diffractogram measured using K α Cu radiation (Bocherens, 1992). When the apatite is more crystalline, the separation between the peaks is bigger and the value of crystallinity index $CI = \Sigma\{H[202], H[300], H[112]\} / H[211] = (a+b+c)/h$ increases (Person *et al.*, 1995).

The mean size of the crystalline domains of bone carbonate hydroxylapatite crystals are measured employing Scherrer's equation: $CS = K\lambda / B \cos\theta$ (Klug and Alexander, 1974) where K is a dimensionless shape factor, with a value close to unity; λ is the X-ray wavelength; B is the line broadening at half the maximum intensity (after subtracting the instrumental line broadening), in radians.

A simple empirical equation to calculate the content of CO₂ proposed by O'Brien *et al.* (1990) were used. This equation is based in the separation of the powder diffraction pattern of the 211 and 112 lines and it is expressed as: $\text{wt \%CO}_2 = 17.335 - [615.524 * (d_{211} - d_{112})]$.

RESULTS

Thin sections and SEM/EDS

Thin sections of cortical and cancellous (trabecular) bone showed the typical birefringence of phosphate minerals, such as apatite.

Cortical bone

The cortical bone retained the normal micromorphology of mammal bones; Haversian system, osteons with the concentric lamellae growing around the Haversian canals and osteocyte lacunae were observed as well as the interstitial lamellae (Figure 6); bone reabsorption can be seen in Figure 7b-1 (Lamm, 2007).

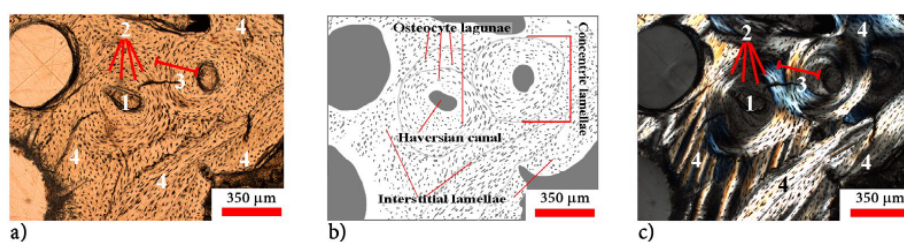


FIGURE 6

a) Thin section of cortical bone under ppl; b) Diagram of cortical bone; c) Same section of cortical bone under xpl. In both micrographs the Haversian system (osteons) is seen: concentric lamellae (3) growing around the Haversian canals (1), osteocyte lacunae (2) are also visible as well as the interstitial lamellae (4).

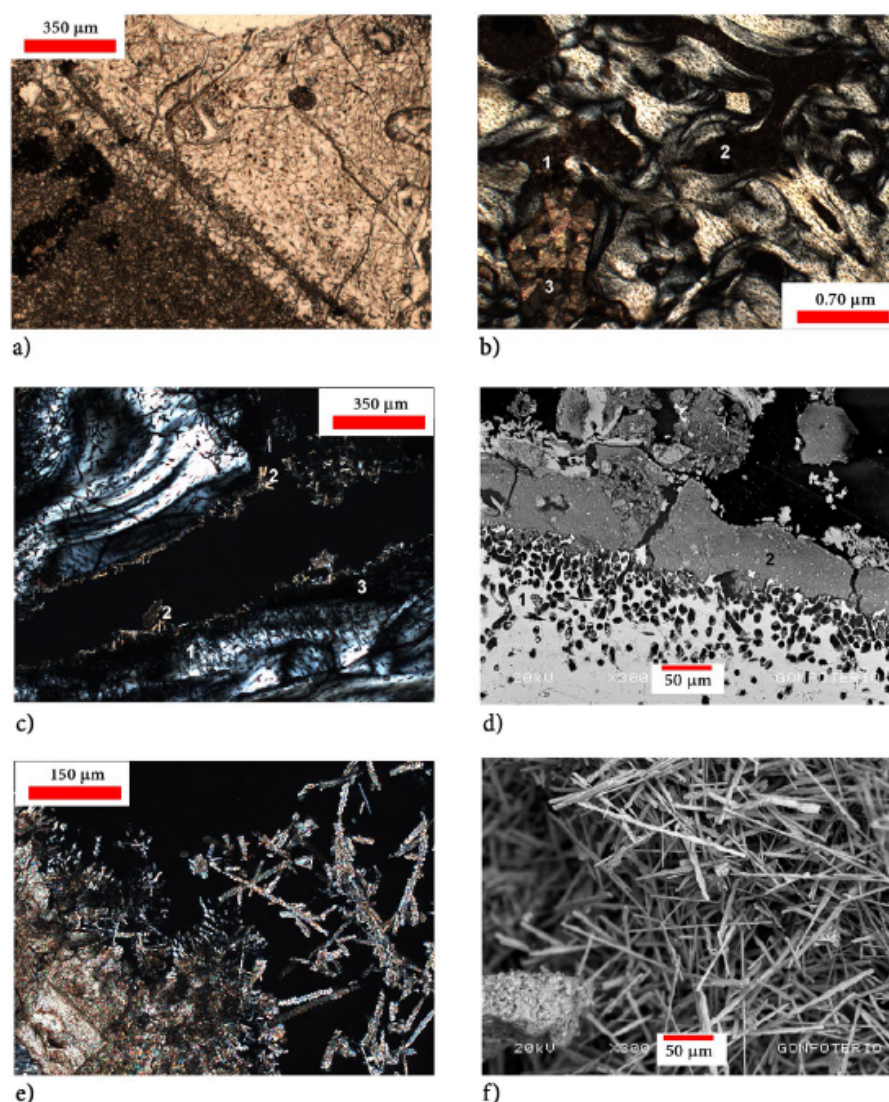


FIGURE 7

Bone alterations. a) Fissures and fractures, only the wider fractures are filled; thin section (ppl). b) Thin section of cortical bone (xpl), 1) reabsorption of the bone visible in Haversian canals that have lost their oval shape, 2) canals filled with micrite, 3) canals filled partially with sparite. c) Thin section of the surface of cortical bone (xpl), 1) tunneling with a rounded end, 2) lublinites in surface, 3) micritized bone, the material have lost their birefringence. d) SEM micrograph of thin section of cortical bone, 1) tunneling with rounded ends, 2) layer with different composition of the bone, not all the tunnels are filled. e) lublinites, thin section (xpl). f) Lublinites, SEM micrograph.

Nevertheless, there were a number of alterations within the cortical bone structure. Firstly mechanical damage was shown in a myriad of fissures all along the cortical bone with no preferential orientation; only the widest and most superficial ones had a micritic filling (Figure 7a).

The canals were, in general, filled with micrite (Figure 7b-2), and in few occasions with sparite (Figure 7b-3); the micritic filling exhibited stains of iron and manganese oxides.

The surfaces of the cortical bone fragments showed dissolution features such as tunnels with a rounded end (Figure 7c1, 7d-1); in an early state they were semi perpendicular to the surface and in a more developed state, they formed a network within a few tens of microns of the bone surface but sometimes within hundred

microns. These cavities were interpreted as the result of selective enhanced dissolution at the areas with weaker/imperfect crystalline structure of phosphate fibers.

Thin sections showed a material with no birefringence in the surface area of the sampled bones composed mainly of Si, Al, O, Mg, Fe, and K (Figure 7c-3 and Figure 8), over this layer a micritic one can be found. At the surfaces that exhibited the tunneling, but not the layer of minerals without birefringence, the voids have been filled with micrite.

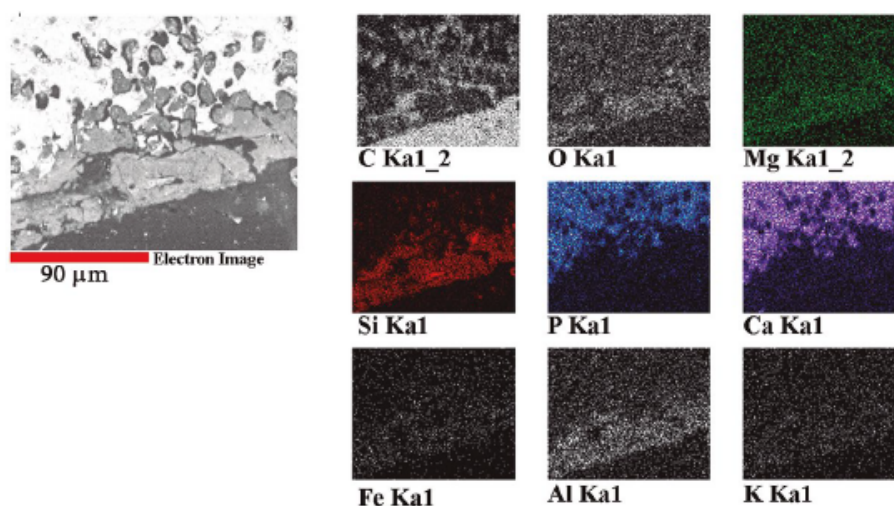


FIGURE 8

SEM image and mapping elements. It can be seen that the filling and layer over the bone is composed mainly of silicon, aluminium, and some oxygen, magnesium, iron and, potassium.

These areas of dissolved bone with outer layers of calcium carbonate had rather similar morphology to that of lublinites -needle fiber calcite (Figure 7c-2). This lublinites is formed by polycrystalline chains, composed of calcite rhombohedron joined together with a P3 Morphology of Verrechia and Verrechia (1994) (Figure 7e and 7f).

Cancellous bone

The morphology of the cancellous bone was also conserved. Parallel lamellae with a plywood structure was visible despite the great amount of fissures and fractures in this type of bone (Figure 9a).

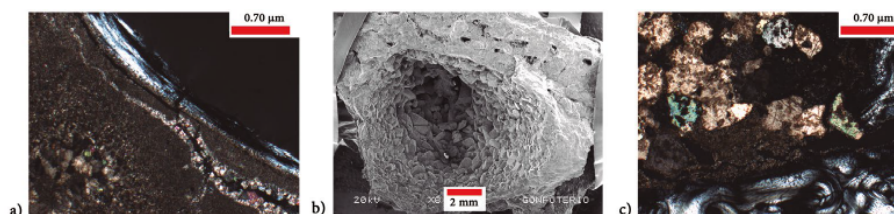


FIGURE 9

- a) Trabecular bone with parallel lamellae and several fissures, the filling consists in a micritic layer with sparitic recrystallization in fractures and other areas, thin sections (xpl). b) SEM image of the crystalline sparitic layer in spaces not completely filled. c) Bone with layer without birefringence, layer of micrite and then sparitic cement, thin section (xpl).

The voids between the walls of the trabecula were filled with specific layers, first a micritic one deposited directly over the surface of the bone (see bottom of figure 9c), then a sparitic layer with big and defined crystals that is the last one (Figure 9b).

When the spaces were filled, after the sparitic layer there was another micritic filling (Figure 9a) and a sandy mixture of minerals and volcanic rocks cemented with micrite or sparite. The minerals in the sand were identified as plagioclase, bone fragments, sparite with the signs of etching, amphiboles, quartz and volcanic glass (Figure 10); we also found a great amount of clays and phytoliths.

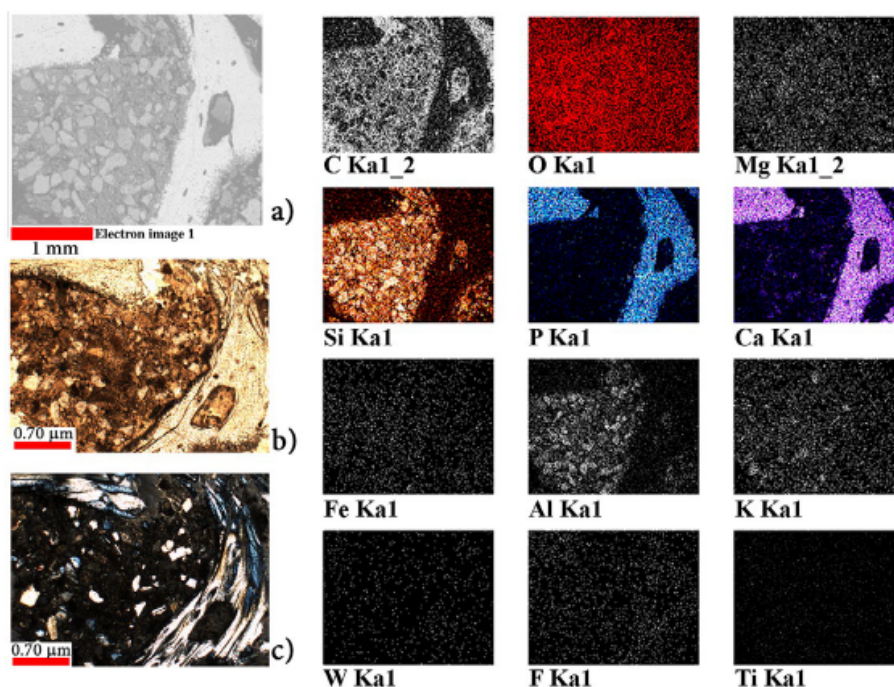


FIGURE 10

Volcanic sand and its elemental composition. a) SEM image of the thin section with a fragment of bone and sand filling. b) Same area in thin section (ppl). c) Same area with cross-polarized light (xpl). At right, we can see the elemental mapping; the volcanic sand fraction is composed mainly by silicium, aluminium and oxygen, with other elements such as magnesium, calcium, iron, potassium.

When the micritic filling had fractures or pores with a large surface, the spaces were filled with sparite.

X ray diffraction

For all the 10 measured bone samples (three aliquots of every type and one duplicate) the peak positions and Miller indexes corresponded to the carbonate-apatite card ICDD 01-073-9696 and general formula: $\text{Ca}_5[(\text{PO}_4)_{3-x}(\text{CO}_3\text{F})_x](\text{F},\text{OH},\text{Cl})$. This structure was used in all the samples in which the Rietveld refinement has been applied. The quantitative results of . and . cell parameters, crystallinity index (CI-DRX), crystallite size (CZ) and wt%CO₂, can be seen in Table 1 and Figures 11, 12 and 13. The *rw*p agreement index for the adjustment of all the samples ranges from 4.8 to 7.0 %.

TABLE 1

XRD values for all the apatite samples: *a* and *c* cell parameters (including error) obtained by Rietveld refinement (*rwp*: agreement index for the adjustment), calculated wt% CO₂ (O'Brien et al. 1992), crystallinity index (CI), and crystal size (CZ, by the Scherrer Equation). DgoApa: mineral apatite from Durango; Cow Bone: fresh bone from a cow.

Sample	<i>a</i> (Å)	<i>c</i> (Å)	wt % CO ₂	CI	CZ (nm)	<i>rwp</i> (%)
1.1	9.3561(7)	6.8989(5)	2.9	0.458	410	6.37
1.2	9.3555(8)	6.8975(5)	3.17	0.434	369	6.42
1.3	9.3577(7)	6.8976(5)	2.95	0.44	463	5.94
2.1	9.3546(7)	6.8973(4)	3.21	0.405	554	6.02
2.2	9.3541(8)	6.8967(5)	3.2	0.392	414	5.48
2.3	9.3568(7)	6.8975(4)	3.21	0.425	433	5.88
3.1	9.3471(3)	6.8958(3)	7.9	0.375	412	5.63
3.2	9.344(3)	6.896(4)	7.9	0.375	482	6.07
3.3	9.3504(6)	6.8959(4)	3.5	0.372	239	6.33
DgoApa	9.3925(4)	6.8831(2)	0	1.25	1568	4.86
CowBone	9.424(1)	6.8922(9)	2.8	0.1	88	7.01

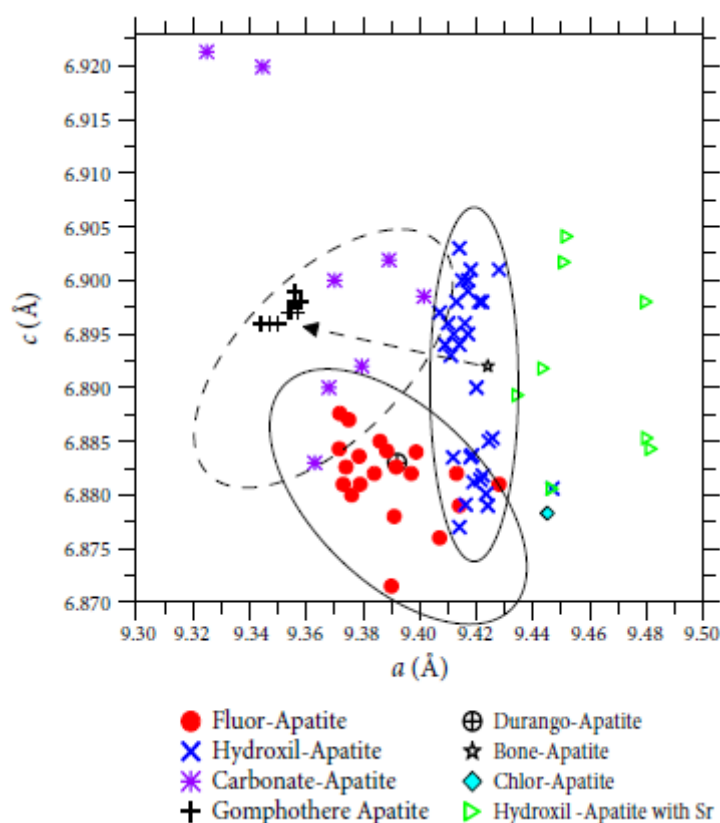


FIGURE 11

a (Å) and c (Å) lattice parameters calculate by Rietveld refinement of the apatite identified in the gomphotere cortical bone fragments (samples 1, 2 and 3) in comparison with a (Å) and c (Å) lattice parameters of the principal different apatite types reported in the literature. The cell parameters of the apatite used for comparative purposes were taken from the international databases of the ICSD (Inorganic Crystal Structure Database) and ICDD (International Center for Diffraction Data). The arrow indicates the process of post-mortem mineralogical transformation. Typical the fluorapatite is the preserved mineral in fossil bones (Wopenka et al 2005) and fluorine proceeds from the interaction between water-soil and sediments.

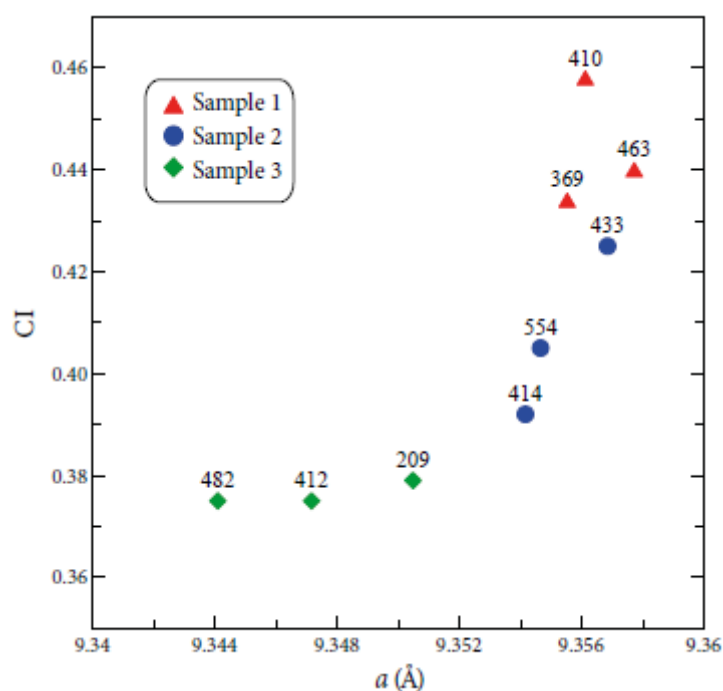


FIGURE 12

Relationship between the Crystallinity Index (CI; Person et al., 1995) and the a (Å) cell parameter for the different aliquots of the three samples of cortical bone. The labels correspond to the crystallite size. It can be observed that number 1 is the bone with the highest crystallinity and the best preserved. Bone number 3 (which probably was the most porous at origin) is the one that mineralogically was more transformed (lower value of a (Å) cell parameter) but has a lower crystallinity index and is slightly less preserved.

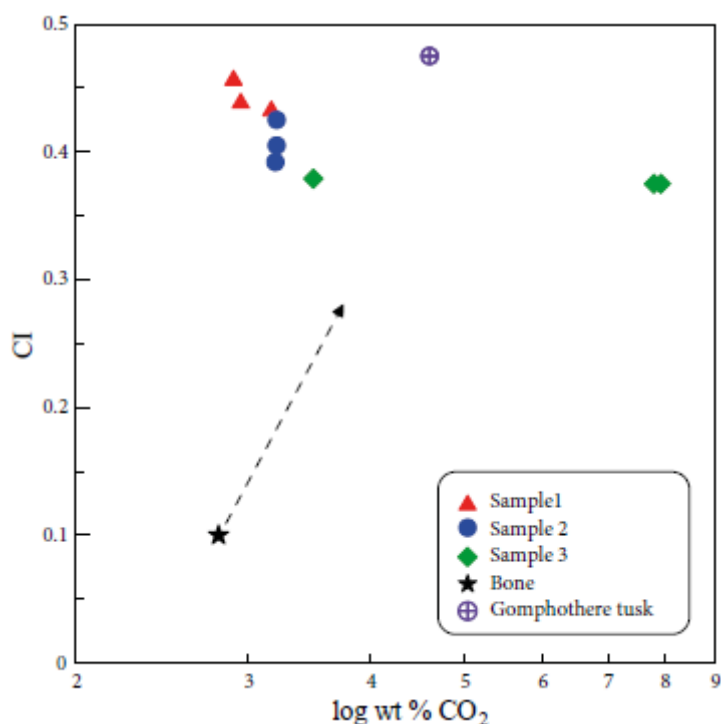


FIGURE 13

Relationship between the Crystallinity Index (CI) and the CO₂ amount (O'Brien et al., 1990) for the different aliquots of the three samples of cortical bone and actual bone. It can be observed that number 1 is the bone with the highest crystallinity index and the lowest CO₂ content. The arrow indicates the general process of post-mortem mineralogical transformation.

During the Rietveld refinement for two samples of group 3, 0.9 % and 1.1 % of smectite clay minerals and less than 1% of quartz (in 1 sample) were found. These features are indicative of detrital minerals filling porosity and cavities. The sediments were composed of calcite, quartz, plagioclase and kaolinitic and smectitic clay minerals.

The clay mineralogy (<2µm) of sediments was composed predominantly of smectite with a small presence of kaolinite. The smectite was confirmed by a strong (001) reflection at about 14 Å in air-dried condition that shifted to about 17 Å in ethylene glycol treated samples and collapsed to 10 Å after heating at 550 °C. The presence of pure smectite (without mixed layers) demonstrates that the diagenetic process that affected the fossil bones was incipient and always bellow 100 °C because this clays begins to transform over these temperatures.

There is a clear correlation between “crystallinity index” and . cell parameter as well as between “crystallite index” and CO₂ content in the carbonate hydroxylapatite (Figures 12 and 13). Samples crystallite size ranged from 370 to 555 nm (with the exception of sample 3.2) in the range reported as the largest dimensions for bones (Glimcher, 1984) confirming the preservation of bone, but smaller than the parameter calculates for geological Cerro del Mercado apatite (1,568nm). For all the analyzed samples, . lattice parameter variations ranged (9.344 Å to 9.357 Å) exceed the variations of c parameter (6.896 Å -6.899 Å).

The aliquots of sample number 3 have small . cell parameter values, more amount of CO₂ and smaller crystallinity index.

DISCUSSION

The transformation in the bone minerals has proceeded in such manner that the paleohistological structure is well preserved. Hubert *et al.* (1996) proposed that when collagen is removed from the structure the bone does not turn into a disorganized mass but the apatite grows on crystal seed preserving the orientation of original crystals and thus preserving micron scale features. Apatite growth stops when the pores and surface of the bone is sealed, in this case, with the calcite layers and clay coatings, and calcite and detrital minerals infillings. The biogenic hydroxylapatite reacted easily with soils and sediments because of its small size and large surface area.

Despite the well-preserved microstructure of the bone, changes in the mineral lattice has been noted on the XRD analysis: the mineralized bone has changed from bioapatite (carbonated form of a non-stoichiometric hydroxylapatite) to carbonate-apatite (*e.g.* card ICDD 01-073-9696) and exhibits higher “crystallinity index”, -smaller a cell parameter than the bioapatite phase characteristic of bones *in vivo*-. Person *et al.* (1995) established that the increase in crystallinity in fossil bones is more related to taphonomic conditions than to the geological age, and that the same variations of crystallinity are found in archaeological and in paleontological bones, which suggests that the increase in crystallinity occurs in the earliest stages of diagenesis.

This mineral transformation can be described as a dissolution-recrystallization process related with decrease in the hydroxyl-ion content (Newesly, 1989). The transformation that preserves the external appearance of the bone was described as a stepwise (isomorphous exchange) non equilibrium transition (Lucas and Prevot, 1991) which is more efficient in porous bone like the one observed in our sample 3. Smaller a cell values in sample 3 were related with further transformation of biogenic hydroxylapatite. The end member of this transformation in a complete fossilization process would be an almost pure fluorapatite as observed in Figure 11 -taken from ICSD (Inorganic Crystal Structure Database) and ICDD (International Center for Diffraction Data) XRD database for a comparative purpose-.

This reduction of the . lattice parameter can be observed (Figure 11) for all the analyzed samples by comparison with power diffraction files (ICSD and ICDD) of pure hydroxylapatite.

Apatite has a hexagonal (P63/m) structure ($a = 9.4\text{--}9.6$, $c = 6.8$ to 6.9 Å) with two metal cation sites, a tetrahedral (T) site (PO_4^{3-}) and an anion column where it is possible to find OH^- , F^- and Cl^- . The carbonate ion can substitute two places in the mineral structure: it can be found at anion column or can substitute phosphate ion at the T site (Hughes and Rakovan, 2015; Wopenka and Pasteris, 2005).

The carbonate in the apatite lattice structure could proceed from neoformed calcite -the main filling mineral in the vascular system of bones and in the voids of the trabecular bone. The calcitic permineralization of bone by micritic precipitation is relatively frequent and it is an usual phenomenon in the fossil record (Fernandez Lopez, 1999); field observations in archaeological sites show that bones are generally preserved when they co-exist with calcite in the sediments (Berna *et al.*, 2004).

The carbonate-apatite mineral phase in the skull bone has a negative correlation between “crystallinity index” and CO_2 amount and a cell parameter (Figures 12 and 13). Carbonate has the effect of making the apatite less crystalline (*e.g.* Nemliher *et al.*, 2004). In the studied samples it can be prove this as an indirect relationship between CO_2 content and “crystallinity index” (Figure 13). Sample 3 is enriched in CO_2 and because of that it is less crystalline and has a smaller crystallite size value. Carbonate in apatite constrains the crystallinity to nanometric scale and gomfotherium fragments have quite big size values of crystallite. Therefore, we suggested a partial isomorphous exchange of carbonate by fluorine in apatite as a result of the diagenetic processes.

Higher crystallinity and stability in the samples with low content of CO_3^{2-} , is related to the fact that Ca-PO_4^{3-} bond is stronger than the Ca-CO_3^{2-} bond, determining higher solubility of carbonate-containing

apatite compared to carbonate-free apatite, thus making the carbonated apatite more susceptible to acid dissolution (Wopenka and Pasteris, 2005).

Secondary features of bone and neoformation of secondary minerals

The presence of the tunneling in the surface of the bone suggests bioerosion due to fungus or cyanobacteria attacks. The term bioerosion is applied to the characteristic destruction of histology, accompanied by an increase of porosity caused by microorganisms in the depositional environment (Turner-Walker and Jans, 2008).

Trueman and Martill (2002) showed that a great amount of fossil bones are affected by bioerosion despite the previous assumption that fossil bones are unaffected by it. The features found in the bones are described as Type 1 Weld tunneling.

Weld described microbial attack on mineralized tissue in 1864. He found tunnels ranging from 5 to 10 μm and under electron microscopy appeared empty with well-defined calcification walls. Hackett (1981) describes thoroughly the Weld tunneling as remarkably uniform in diameter, always passing away from the surfaces of the cortex and the osteon canals; also, their course is never smooth, straight nor longitudinal between its main features. The tunneling in the samples fulfilled all of these features (see figure 2a and 3b of Turner-Walker and Jans 2008, figure 1 and 2 on Jans, 2008; figure 2a on Trueman and Martill 2002, figure 2 and 3 of Hackett 1981 and compare to Figures 7c y 7d here).

It is not rare to find Weld tunneling in the samples because it is by far the most common type of bioerosion (Trueman and Martill, 2002) and it is usually found in animal bones (Jans *et al.*, 2004). Fungal attack is the recognized cause for this type of tunneling but also has remarkable similarities to those made by endolithic filamentous cyanobacteria; despite being phototrophic, the cyanobacteria can thrive in environments with extremely low light levels (Turner-Walker and Jans, 2008).

Jans (2008) showed that the fungal hyphae contained minerals in solution within them, while the tunnels do not have any redeposited mineral; this indicates that the dissolved apatite is transported out of the bone as shown in the micrographs of Figures 7c and 7d. It is not clear if fungi dissolve the bone to harvest nutrients or as a medium; as well the tunneling caused by cyanobacteria does not show redeposited minerals.

Cyanobacteria depend on light, so the surface of the bone will be exposed to dissolution only until the bone is covered with sediments. On the other hand, fungus can grow and thrive without light and the attack could have taken place in almost every stage before the bone was entirely sealed.

It is not possible to know if the attack was due to cyanobacteria or to fungus so we cannot establish if the biological attack of the skull was in early times prior to its burial or when it was buried with sediments. However, the probability of strong biogenic alteration of bone after burial seems to be rather low, because the sediments of Layer A have very weak signs of pedogenetic and biological processes other than carbonate migration. Only fragments of tephate rock show features of pedogenesis, regardless of their redeposition and not indicative of soil formation at the site. Therefore, the possibility of bone weathering and phosphate mineral alteration on the surface before incorporation into colluvial stratum is rather high.

In addition, the process of bioerosion is suggested by the presence of needle fiber calcite in the micritic layer directly over the surface of the bone. This mineral suggests weathering due to direct or indirect biological agents and with morphology changes due to physicochemical processes (Verrecchia and Verrecchia, 1994; Becz-Deak *et al.*, 1997; Khokhlova *et al.*, 2001; Bajnoczi and Kovacs-Kis, 2006; Durand *et al.*, 2010; Straulino *et al.*, 2013).

The neoformation of various secondary minerals within the pore space of the bone, as well as the transformation of the primary mineral components of bone, are related to a set of processes. These processes comprise the transformation of the primary structure, composition of bone tissue, deposition of minerals

and the neoformation of various secondary minerals in the surface and porous structure of the cortical bone and in the voids of the trabecular bone.

A relative chronology and succession of the processes according to the relative position of minerals on microscale is proposed below:

Collagens loss. Isotopes analysis showed N value of zero and ^{13}C value of -26.3, which indicates that the carbon is from an external source (Cienfuegos-Alvarado and Perez-Crespo pers. com.)

Heterogeneous mineral transformation of the bone hydroxylapatite.

Partial dissolution of phosphates in the cortical bone surface as shown in figures 7c and 7d that seems to be related with microorganisms.

Precipitation of clay suspensions. The layer immediately over the bone composed of Si, Al, O, Mg, was identified as a clay with no birefringence; blocks of oriented clays in the trabecular bone were identified. With XRD, the clay was identified as smectite and some kaolinite. It is also important to point out that the presence of pure smectite confirms that the diagenesis was minimum.

Precipitation and crystallization of secondary calcite in the bone voids: a) quick precipitation of micrite, b) slower crystallization of sparite in fissures and voids. Precipitation of fine layer of micrite in the bone surface and its transformations to needle fiber calcite (lublinite).

Deposition of sand particles such as volcanic rocks, plagioclase, bone fragments, etched sparite, amphiboles, quartz and volcanic glass. Secondary calcite precipitation at the same time.

The trabecular bone exhibits a bad state of preservation mainly due to physical causes (it is substantially fragmented and fissured). The presence of smectite in the voids of trabecular bone (see section 5.2) could lead to the expansion of this clay causing stress in the bone and consecutively the release of this stress by the fragmentation of the bone.

Types and chronology of these processes in the Cuenca Alta of Tlaxcala could be related to the pedogenetic processes that developed in the sediments; however, the sediments of the site are currently in analysis.

On the other hand, similar progressions have been studied in detail in the key sections of the Tlaxcalan localities of Tlalpan and Mamut (Sedov *et al.*, 2009) where it was shown that weathering, clay neoformation and clay illuviation took place in the late Pleistocene (MIS3-2), whereas neoformation of carbonates occurred predominantly in the Holocene. It is possible that such succession is valid also for the studied mineralization processes.

Implications in different areas

The paleoclimatic implications point out that the weathering, microbial transformations, and clay illuviation are indicators of humid climate; while carbonate depositions evidence aridization.

Secondary carbonates and layers of clays seal the bone surface and thus prevent further deterioration promoting the preservation of the bone.

Implication for the geochemical records implies that secondary carbonates not only precipitate on the bone and teeth surface, but also penetrate pores and fractures; thus they could influence chemical and isotopic composition of the bone samples altering the geochemical record of past environment.

CONCLUSIONS

Bone paleohistological structure is well preserved; however, the cortical bone is more preserved than the cancellous bone because the clays that fill the trabecula macroporosity generated internal stresses that fragmented the bone walls.

Nevertheless, the bone composition changed; there is no collagen and there is exogenous C; the mineral part of the bone also was transformed from carbonate non-stoichiometric hydroxylapatite to more crystalline apatite- probably fluorine-carbonate apatite. These changes imply a more stable and less soluble mineral than that of the original bone.

The exogenous carbon detected with isotopical analysis and the carbonate phase in the carbonate-apatite could proceed from the calcite, the main filling mineral. The permineralization of bone by calcite is frequent and enhance the bone preservation.

Biological activity was detected by dissolution and structure modification of the bone surface and the transformation of micrite to needle fiber calcite.

With the calcite infilling we also observed clays, pure smectite and kaolinite, which could be the first minerals intruding the bone macroporosity and deposited in the bone surface. Furthermore, we found sand filling composed of quartz, feldspars, amphiboles and volcanic glass, with a clear different origin if compared with the clays and the calcite.

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