

A Pompeii-like preservation for *Dosinia ponderosa* (Gray, 1838): two volcanic ash casts from the East Pisco Basin (Miocene, Peru)

Giulia Bosio

G. Bosio, Dipartimento di Scienze dell' Ambiente e della Terra, Università degli Studi di Milano-Bicocca, I-20126 Milano, Italy; giulia.bosio.giulia@gmail.com

KEY WORDS - *Taphonomy, tephra, fossil-diagenesis, carbonate dissolution, distal ashfall, pyroclastics.*

ABSTRACT - *In the East Pisco Basin, in Peru, a highly unusual example of Pompeii-like preservation has been discovered. At the locality of Cerros la Mama y la Hija, along the western side of the lower Ica River Valley, south of Ocucaje, two bivalve specimens were found preserved as volcanic ash casts. As is the case for most fossil invertebrates from the Pisco Formation, the original carbonate shell is not preserved, but these specimens are nonetheless exquisitely preserved, even featuring taxonomically informative characters of the shell exterior that allow for their assignment to the locally common species *Dosinia ponderosa* (Gray, 1838). Both casts are mainly constituted by fine-grained volcanic glass shards and biotite crystals. The high abundance of glass shards and the absence of non-volcaniclastic material such as biogenic or terrigenous particles indicates that the casts are the product of primary deposition of a volcanic ash layer on the seafloor. These specimens testify to an intriguing taphonomic history that features burial underneath a distal volcanic ashfall. During the Late Miocene, pyroclastic ashfalls coming from the Central Andes were indeed frequent in the East Pisco Basin, allowing this exceptional type of fossil preservation.*

INTRODUCTION

The Peruvian East Pisco Basin is globally known for its paleontological content. The Miocene Pisco Formation (Fm) hosts one of the major marine vertebrate Fossil-Lagerstätte of the Cenozoic Era, which includes fossil cetaceans, pinnipeds, fishes, seabirds and marine reptiles (e.g., Lambert et al., 2010; Esperante et al., 2015; Bosio et al., 2021b; Collareta et al., 2021; Bianucci & Collareta, 2022; Bianucci et al., 2023). Abundant and diversified fossil assemblages of marine invertebrates, comprising mollusks, acorn barnacles, cirratulids, echinids, brachyuran decapods, rare bryozoans, and benthic foraminifera are also found in this basin (e.g., Lisson, 1925; Rivera, 1957; Alleman, 1978; DeVries, 1988; DeVries & Frassinetti, 2003; Coletti et al., 2019a, b; Collareta et al., 2019, 2023; Villaseca et al., 2020; Bosio et al., 2021a; Koči et al., 2021; Sanfilippo et al., 2021).

In the Pisco Fm, paleontologists can find a variety of fossil preservation styles. Vertebrate remains are found partially or totally enclosed in dolomite nodules or, more frequently, they are characterized by extensive bone phosphatization. Therefore, there are two main mineralization processes which allow bone fossilization: the dolomite precipitation and the apatite dissolution-recrystallization (Gariboldi et al., 2015; Gioncada et al., 2018a; Bosio et al., 2021b, c). Different preservation modes are also recognized for fossil invertebrates in the three depositional sequences of the Pisco Fm (Di Celma et al., 2017). In particular, in the Middle Miocene strata, fossil preservation consists of calcite internal molds, with both pristine and recrystallized calcite shells (Di Celma et al., 2017; Bosio et al., 2020b). In the Upper Miocene units, fossil invertebrates are usually preserved as dolomite internal, external and compound molds, and/or as gypsum/anhydrite molds, recrystallized shells or casts. In these strata, the original mineralogy of the carbonate shells

is not preserved (Di Celma et al., 2017; Gioncada et al., 2018b; Bosio et al., 2021a; Koči et al., 2021; Sanfilippo et al., 2021).

Here, a highly unusual preservation style is reported for the first time in the East Pisco Basin for fossil invertebrates. Two fossil specimens of *Dosinia ponderosa* (Gray, 1838) were found preserved as casts of volcaniclastic material in a surrounding sandy sediment. These volcanic ash casts represent the only reported find of this type of preservation in the Peruvian fossil record and one of the few cases in the world.

GEOLOGICAL AND PALEONTOLOGICAL FRAMEWORK

Tectonic and stratigraphic setting

On the South American plate, a NW-SE series of Cenozoic marine basins runs along the Peruvian coast (Thornburg & Kulm, 1981). Among these forearc basins, the East Pisco Basin mainly extends between the towns of Pisco in the north and Nazca in the south and is mostly exposed onshore (Fig. 1). Sediment deposition in this marine basin has been active from the Eocene to the Pliocene, when the most recent exhumation of the marine sediments started following the subduction of the Nazca Ridge (Hsu, 1992). These sediments are organized in the following formations, from the oldest to the youngest: the Caballas Fm, the Paracas Fm, the Otuma Fm, the Chilcatay Fm and the Pisco Fm (Dunbar et al., 1990; DeVries, 2017; DeVries et al., 2017; DeVries & Jud, 2018; Di Celma et al., 2022). Along the western side of the Ica River Valley, south of the village of Ocucaje, the Pisco Fm is Middle to Late Miocene in age and comprises three depositional sequences, namely P0, P1 and P2 in stratigraphic ascending order, bounded by stratigraphic unconformities, PE0.0, PE0.1 and PE0.2, respectively (Di

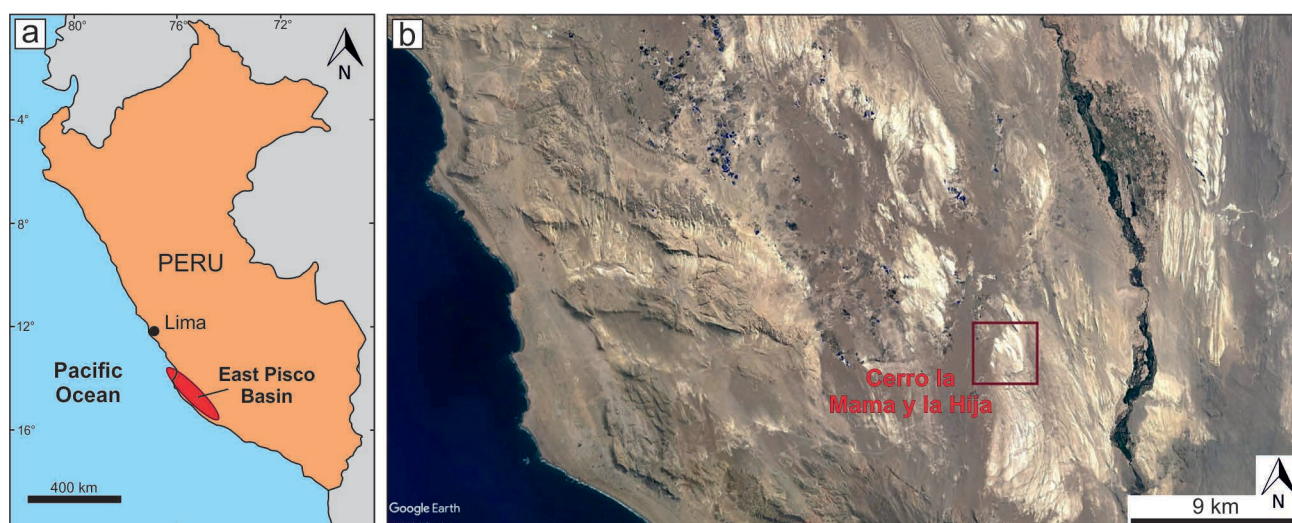


Fig. 1 - (color online) Geographic context. a) Geographic location of the East Pisco Basin along the Peruvian coast. b) Satellite image of the locality of Cerros la Mama y la Hija (Ica Desert, Peru). Based on Google Earth image.

Celma et al., 2017). Each stratigraphic unit is composed of sandstones at the base and siltstones or diatomaceous siltstones at the top, representing a transgressive cycle (Di Celma et al., 2017). The P0 sequence is dated between 14.7 and 12.6 Ma (Bosio et al., 2020b, 2022), the P1 sequence between 9.5 and 8.6 Ma, and the P2 sequence between 8.4 and ≥ 6.71 Ma (Bosio et al., 2020c). All the stratigraphic units record long-lasting, explosive volcanic activity. Numerous volcanic ash layers are found within the marine sediments, testifying to pyroclastic fallout in the area, coming from both the Western and Eastern Cordillera (Bosio et al., 2019, 2020a, b).

Invertebrate paleontology

In the East Pisco Basin, fossil invertebrates can be found throughout the sedimentary succession, from the middle Eocene Los Choros Member of the Paracas Fm to the Upper Miocene strata of the Pisco Fm, with a very wide range of fossil preservation types. Since the 1980s, much attention has been given to fossil mollusk assemblages from the East Pisco Basin, including descriptions of new species (e.g., Muizon & DeVries, 1985; DeVries, 1988, 1997, 2001, 2007, 2016, 2019; DeVries & Frassinetti, 2003; DeVries et al., 2006; Bosio et al., 2021a; Sanfilippo et al., 2021). The Pisco Fm hosts abundant molluscan remains, usually concentrated in mollusk-rich layers with low biodiversity and high dominance (Bosio et al., 2021a) and rarely associated with vertebrate remains (Bosio et al., 2021b).

Along the western side of the Ica River Valley, in the P0 sequence, the Middle Miocene molluscan fauna suggests warm-water conditions, as also indicated by the vertebrate fauna (DeVries & Frassinetti, 2003; Bosio et al., 2020b; Collareta et al., 2021). An important invertebrate faunal change occurred through the PE0.1 unconformity due the appearance of new species that show affinities with the Miocene Panamic fauna rather than with that of the Lower to Middle Miocene Navidad Fm of Chile, suggesting cooler conditions (DeVries & Frassinetti, 2003; Nielsen & Glodny, 2009; Di Celma et al., 2017).

Regarding bivalves, specimens of *Dosinia ponderosa* are found in all the three depositional sequences of the Pisco Fm (Di Celma et al., 2017). Today, this species is typical of intertidal zones down to depths of 60 m (Coan & Valentich-Scott, 2012), suggesting a shallow-water marine paleoenvironment at the locality of Cerros la Mama y la Hija during the Late Miocene. Even if the molluscan fauna of the Upper Miocene Pisco Fm suggests cooler conditions than Lower and Middle Miocene fauna, showing affinities with the Panamic fauna (DeVries & Frassinetti, 2003; Di Celma et al., 2017), *Dosinia ponderosa* is usually associated with warm waters and its disappearance from the southern Peruvian coast could be attributed to the onset of Pleistocene cooling (DeVries, 1986; DeVries & Frassinetti, 2003) and/or to the upwelling conditions due to the strengthening of the Humboldt Current that started in the Late Miocene and continued until today (Amiot et al., 2008; Bosio et al., 2020b; Collareta et al., 2021; Kiel et al., 2023).

MATERIAL AND METHODS

Study area and sample collection

The P1 sequence of the Pisco Fm displays a maximum thickness in its southern outcrops, at the locality Cerros la Mama y la Hija ($14^{\circ}36' S$; $75^{\circ}41' W$) (Figs 1b, 2). Here, the P1 strata consist of sandstones at the base, and the rest of the succession is composed of diatomaceous siltstones alternating with dolomite nodules and volcanic ash layers (Fig. 2a). At the top, a stratigraphic unconformity separates the P1 sequence from the overlying P2 sequence, starting with a lag of authigenic nodules and sandstones (Fig. 2a). At this locality, many fossil vertebrates have been discovered, such as the mysticete specimen shown in Fig. 2c (Esperante & Poma, 2015; Collareta et al., 2021), and several volcanic ash layers have been studied and collected (Fig. 2d). The two mollusk specimens that are studied herein were collected at the base of the hill: MUSM-INV308 from the northern side of these hills

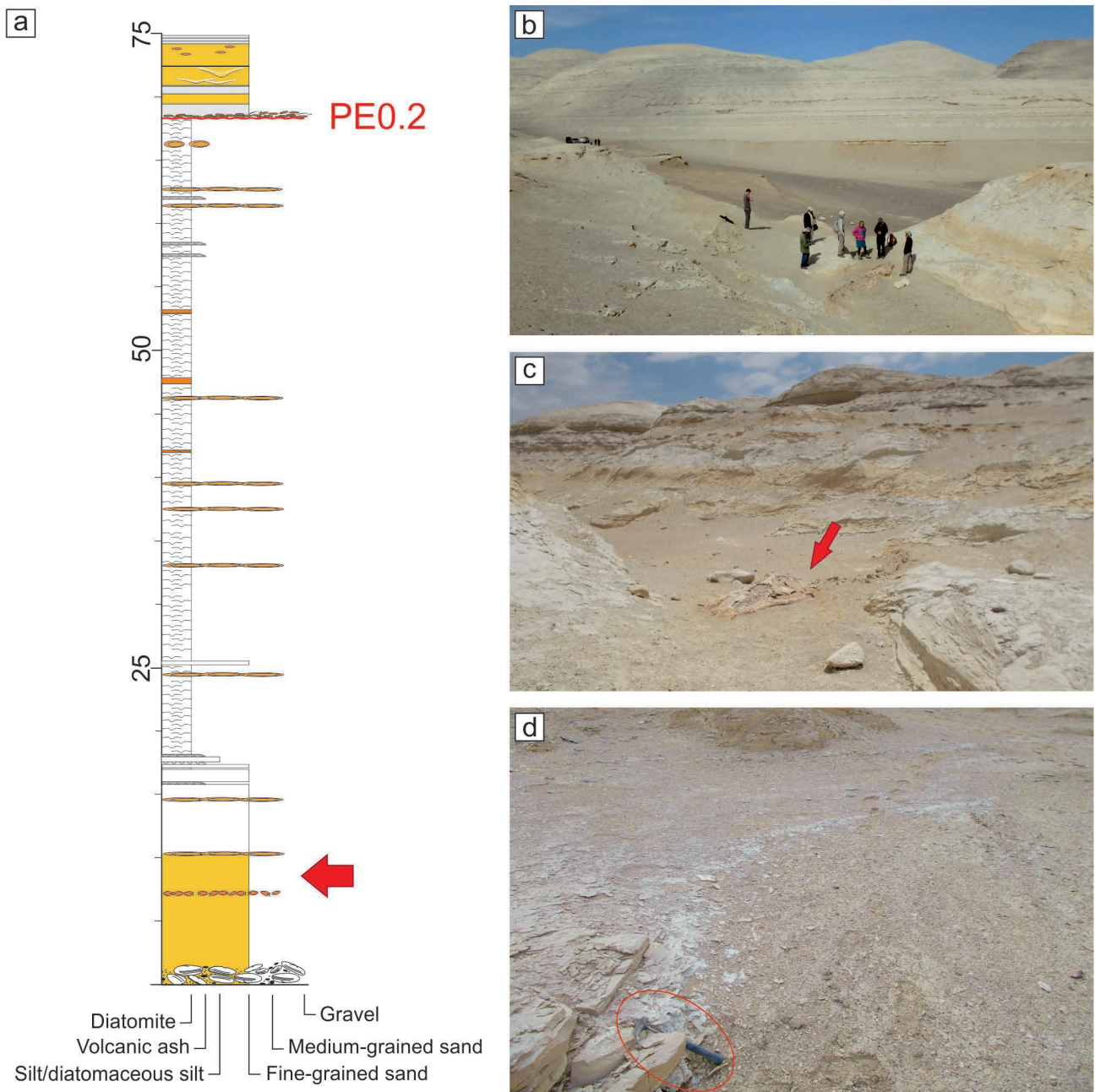


Fig. 2 - (color online) Stratigraphy and field photos. a) Measured stratigraphic section at the locality of Cerros la Mama y la Hija. The red arrow indicates the layer of the collected bivalve samples. b) Sandstones and diatomites of the Pisco succession cropping out at the locality of Cerros la Mama y la Hija. Note that sandstones are the orange sediments in the lower part of the photo, whereas the diatomaceous silstones are whitish in color and exposed at the top of the succession. c) Fossil specimen of a mysticete (red arrow) in the sediments of the P1 sequence at the locality of Cerros la Mama y la Hija. d) A volcanic ash layer (whitish layer) in the sedimentary succession of the P1 sequence at the locality of Cerros la Mama y la Hija. The red circle spotlights the hammer in the left corner.

during fieldwork in June 2017, not in situ ($14^{\circ}35'56.5''$ S; $75^{\circ}40'58.4''$ W), and MUSM-INV309, in situ, from the southern side of the hills during fieldwork in September 2021 ($14^{\circ}36'37.8''$ S; $75^{\circ}40'41.4''$ W). The two specimens were described and deposited in the invertebrate paleontology collection at the Museo de Historia Natural de la Universidad Nacional Mayor de San Marcos (MUSM, Lima, Peru).

Laboratory analyses

The two studied specimens were described in the laboratory and measured with a standard analog caliper.

Two smear slides were prepared from the unconsolidated material forming the casts for investigating the composition of the two bivalve specimens. A small amount of sediment composing the mold was collected from the specimens with a wet toothpick and rubbed on the microscope slide with a drop of distilled water. Slides were positioned on an electric burner at ca. 50°C and dried. They were then covered with a cover glass using a Norland Optical Adhesive and exposed under a UV lamp. The two smear slides were analyzed through an Optika B-500POL optical microscope under transmitted light.

RESULTS

At Cerros la Mama y la Hija, specimen MUSM-INV309 was found in an indurated layer within the sandstones at the base of the succession (Figs 2a, 3a-c). This specimen was preserved in situ but not in life position, with the posterior part pointing to the stratigraphic bottom (Fig. 3a-c). The other specimen, MUSM-INV308, was found ex situ, on the desert floor, far from the outcrop (Fig. 3d). Both the specimens are white-colored fossil casts of uncemented and scarcely consolidated volcanic material consisting of fine-grained glass shards and biotite crystals observable through a hand lens. They are surrounded by a coarser-grained brownish sediment with the grain size of medium sand. The casts are partially eroded or display abrasion marks due to recent wind action. Except for these slightly eroded parts, the fossils are exquisitely preserved, retaining the original shell morphology. The casts of the shells are subcircular in shape, moderately inflated, equivalve and slightly inequilateral (Fig. 4). The lunula is not well-preserved or is covered by sediment, whereas the escutcheon is obscure (Fig. 4). The umbos are pointing anteriorly, indicating that the shell is prosogyrate (Fig. 4). They exhibit an evident sculpture of commarginal

ribs with very narrow interspaces (Fig. 4). The MUSM-INV308 specimen measures 7.22 cm in height, 7.62 cm in length, and 3.20 cm in width; the L/H ratio is 1.05. The MUSM-INV309 specimen measures 7.19 cm in height, 7.82 cm in length, and 2.90 cm in width; the L/H ratio is 1.09. Both the casts have a small portion lacking in length. By comparing shell characteristics and measurements with published data about the tropical West American molluscan fauna (Coan & Valentich-Scott, 2012) and the Pisco Fm molluscan fauna (DeVries, 1986; DeVries & Frassinetti, 2003; Bosio et al., 2021a), the casts are attributed to the species *Dosinia ponderosa* (Gray, 1838). The values of height, length and width and the L/H ratios perfectly overlap those reported by Bosio et al. (2021a) for internal dolomite molds preserved in the P1 strata.

Since the grain size of the poorly consolidated volcanoclastic material is below 2 mm, the bivalve casts are composed of ash (White & Houghton, 2006). Under the optical microscope, ca. 90-95% of the components are volcanic glass shards, whereas the remaining 5-10% is composed of biotite crystals (Fig. 5). Glass shards are mainly highly vesiculated and stretched (Fig. 5). Since the extraneous particles (i.e., biogenic material, rock fragments, iron oxides, clays) represent less than 5%,

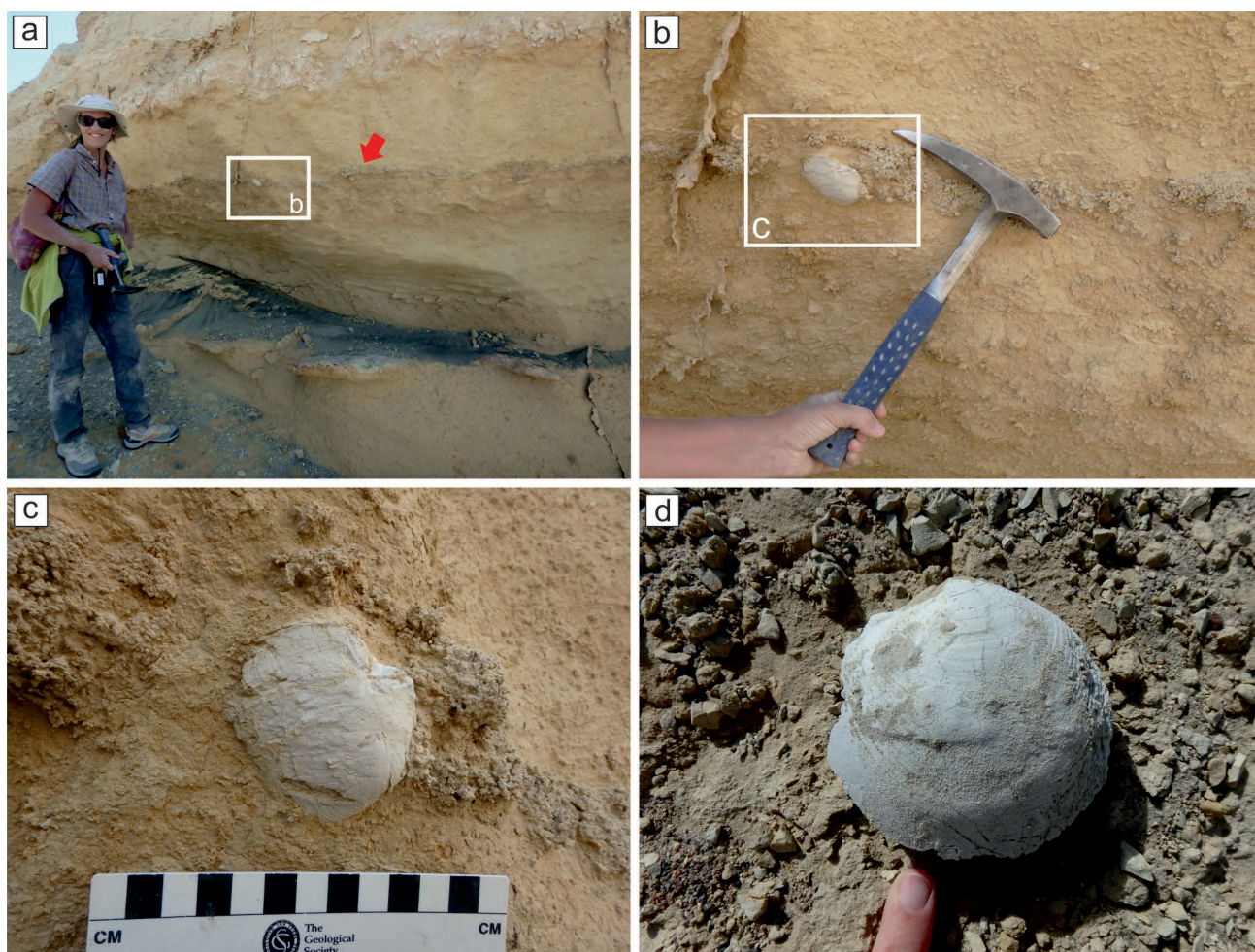


Fig. 3 - (color online) Field photos of the bivalve specimens. a) Field photo of the P1 sandstones cropping out at Cerros la Mama y la Hija where one of the samples was collected in situ. The red arrow indicates the layer of the collected MUSM-INV309 specimen. b) Close-up of the boxed area in Fig. 3a showing the layer with the bivalve MUSM-INV309. c) Close-up of Fig. 3b showing MUSM-INV309 specimen. d) MUSM-INV308 specimen found not in situ at the base of the sandstones of the P1 sequence cropping out at Cerros la Mama y la Hija.



Fig. 4 - (color online) MUSM-INV308 specimen. Fossil cast of *Dosinia ponderosa* (Gray, 1838) from Cerros la Mama y la Hija, Peru. From upper left clockwise: left valve, right valve and dorsal view. Height: 7.22 cm; length: 7.62 cm; width: 3.20 cm. Scale bar corresponds to 2 cm.

the casts' components represent the product of primary deposition of a volcanic ash layer, excluding a reworking (see Griggs et al., 2014; Bosio et al., 2019).

DISCUSSION

Fossil conservation in volcanic ash

There are not many cases of fossils preserved in volcanic ash, but these are usually exquisitely preserved (see Lockley, 1990). It is worth mentioning the archeological site of Pompeii and its external molds formed within the body of pyroclastic materials (e.g., Giacomelli et al., 2003; Deem, 2005). Today, studies on recent ashfall deposits have been carried out to understand taphonomic processes in these particular settings (e.g., Hayward et al., 1989; Heikoop et al., 1996b; Lokier, 2021; Maguire, 2022).

Ashfalls allow the exquisite preservation of fossil traces (e.g., Hay & Leakey, 1982; Baales et al., 2002; Mietto et al., 2003; Nelson et al., 2008; Marriott et al., 2009) and both mineralized and non-mineralized bodies by instantaneously blanketing entire forests and terrestrial plant remains - the "Pompeii effect" (e.g., Wagner, 1989; Scott, 1990; Hayward & Hayward, 1995; Goetcheus & Birks, 2001; Pfefferkorn & Wang, 2007; Opluštil et al., 2009; DiMichele & Falcon-Lang, 2011; Rößler et al., 2012; Locatelli, 2014; Góis-Marques et al., 2019; Wick,

2023) - as well vertebrate fauna in both continental (e.g., Hay, 1986; Famoso & Pagnac, 2011; Antoine et al., 2012; Iurino et al., 2014; Jiang et al., 2014) and marine (Bosio et al., 2021b) settings, and benthic communities on the seafloor (e.g., Heikoop et al., 1996a; Droser et al., 2006; Maguire et al., 2016). One of the most renowned examples of the lattermost case is the Ediacaran fauna, which is characterized by different types of preservation, all sharing the rapid burial by event beds, such as instantaneous deposit of volcanic ash on the seafloor to form an "Ediacaran Pompeii" (Seilacher, 1992; Narbonne, 2005; Droser et al., 2006; Hofmann et al., 2008). The predominant type of preservation is a "conception-style preservation", which consists of epireliefs (positive reliefs) such as casts of ash molds on a surface overlain by a volcanic ash bed (Narbonne, 2005; Bamforth et al., 2008). Fossil preservation turns out to be exceptional also in other volcanic settings such as maar lakes (e.g., Kovács et al., 2020; Grandi et al., 2023)

The extraordinary preservation in volcaniclastic material is due to the rapid burial below a thick and heavy load of ashfall and the early diagenesis of labile minerals, which allows a rapid cementation in the volcanic ash that contribute to preserving organisms (Heikoop et al., 1996a; Narbonne, 2005; Opluštil et al., 2009; DiMichele & Falcon-Lang, 2011). The instantaneous burial rapidly isolates the organisms from the destructive effects of subsequent hydraulic disturbance, scavenging, bioerosion

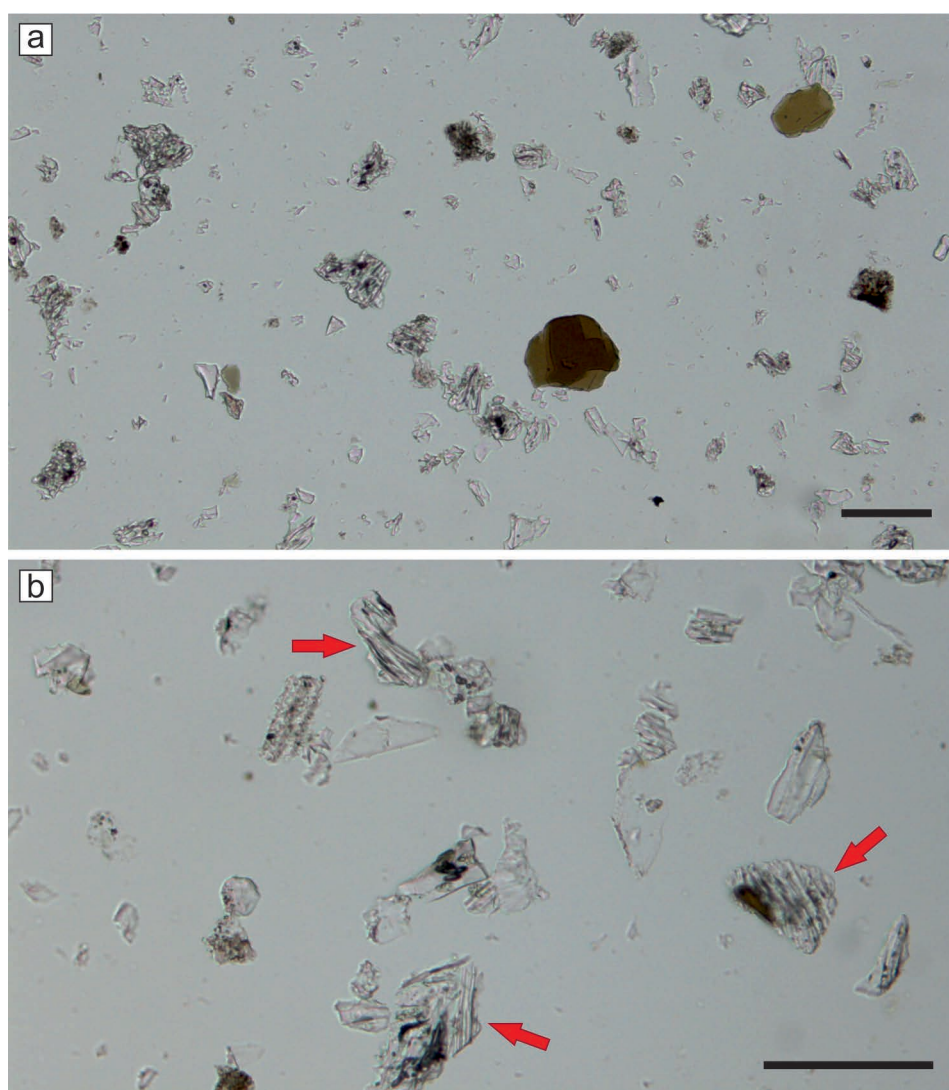


Fig. 5 - (color online) Photomicrographs. a) Highly vesiculated volcanic glass shards (transparent) and biotite crystals (brown color) forming the bivalve casts observed under the optical microscope. b) Detail of the highly vesiculated and stretched volcanic glass shards (red arrows). Scale bars equals 100 μm .

and bioturbation (Heikoop et al., 1996a). Droser et al. (2006) assert that organisms already lying on the substrate are cast with precision, while those transported by the ashfalls are less well molded. The preservation potential for organism details could be also limited by the crystal size of the volcanic grains (Narbonne & Gehling, 2003) and the temperature of the volcanic material (Iurino et al., 2014). In fact, the fine-grained material favors the formation of external molds and the low temperature phreatomagmatic deposits preserve detailed paleontological information that are normally lacking in the fossil record (Iurino et al., 2014). The variety of preservation includes cellular permineralization, also defined “petrification” for plants, adpressions, epireliefs, casts and molds (Rößler, 2021). In some cases, volcanic ash deposits preserved fossils in carbonate concretions (Orr et al., 2000; Briggs et al., 2008).

A taphonomic enigma

The two volcanic casts of *Dosinia ponderosa* (Gray, 1838) discovered at the locality of Cerros Mama y la Hija

tell an intriguing taphonomic story. The exact order of the processes which brought about the formation of ash casts within a sandy sediment remains an enigma.

Dosinia ponderosa is a mobile infaunal suspension feeder, with a compressed shell shape and vertical life position (Di Celma et al., 2002). This species lives in substrates of fine sand to coarse silt characterized by low to high energy (Baqueir, 1979) in a nearshore zone shallower than 60 m (Coan & Valentich-Scott, 2012). During the Late Miocene, *Dosinia ponderosa* specimens lived in the shallow waters of the East Pisco Basin within a sandy sediment with sedimentological features attributable at ca. 5-10 m of depth (see Sanfilippo et al., 2021 for an analog from the basal sandstones of P2). After death, infaunal mollusks tend to stay buried within the sediment, while epifaunal mollusks tend to remain exposed on the sediment-water interface (Lazo, 2004). Therefore, infaunal bivalves such as *Dosinia ponderosa* have a possibly higher fossilization potential and time averaging than the epifaunal bivalves (Lazo, 2004). Because we found the in situ specimen not in life position,

both Pisco specimens would have been slightly rotated by bioturbation or exposed on the seafloor by a storm event without being disarticulated (Fig. 6). On the seafloor, they were soon covered either by a sandy sediment or by a centimeters-thick volcanic ash layer erupted by volcanic centers in the near Central Andes (Bosio et al., 2020a) (Fig. 6). This rapid burial probably prevented the disarticulation of the shells.

In the first hypothesis (see Fig. 6 left), sand surrounding the bivalves started to consolidate, allowing the formation of an external mold in the sediment. At the boundary between the sulphate reduction and methanogenesis zones, pH variations may occur during early diagenesis (Meister et al., 2007, 2011; Xu et al., 2019). A decrease of pH would explain the dissolution of the aragonitic shell, aragonite being unstable and easily dissolved (Cherns et al., 2011). After these processes, two empty external molds remain in the consolidated sediment. Then, an event such as a storm removed the first few centimeters of lesser consolidated sediments, and a small aperture opened at the top of the cavities. Immediately, a volcanic ashfall covered the seafloor filling the cavities. In shallow water settings, volcanic deposits have a very low preservation potential due to the presence of syndepositional disturbances, such as the action of marine currents and storm waves (Carey, 1997; Griggs et al., 2014; Bosio et al., 2019). In this case, a winnowing event occurred, removing the ash layer and leaving intact the two filled cavities on the sediment surface. After winnowing, the two casts were buried in newly deposited sandy sediment and stayed fossilized until the recent exhumation.

In the second scenario (see Fig. 6 right), during burial by the primary deposition of a volcanic ash, the two specimens were filled by the ash. Due to the fine grain size of the ash, winnowing occurred, but the articulated bivalves preserved the compact and stiff ash inside the shell. After the winnowing, the filled shells were buried again by sandy sediment deposited on the seafloor. After a long time, the surrounding sediment started to cement and an external mold of the shells formed. During diagenesis, fluids enriched in silica, probably due to the highly siliceous environment (e.g., volcanic ash, terrigenous sediments, overlying diatomaceous silts) dissolved aragonite of the shells and remobilized the fine-grained ash inside the internal casts, which took on the form of the external mold of the shells. In fact, the release of dissolved silica and cations such as $\text{Fe}^{2+/3+}$, Mn^{4+} , and Ba^{2+} from volcanic glass shards in marine sediments is always accompanied by a strong decrease in alkalinity and pH in the surrounding waters (Kutterolf et al., 2006).

A difficulty with the first scenario is that an unusual series of synchronized processes needed to have occurred. The problem with the second scenario is that in the last process, where the shell dissolved and ash adapted to the external molds, a diminishment of volume should have occurred (and this is not observed).

CONCLUSIONS

Casts of two bivalve specimens collected at the locality of Cerros la Mama y la Hija in the East Pisco Basin (Peru) exhibit exquisitely preserved features of

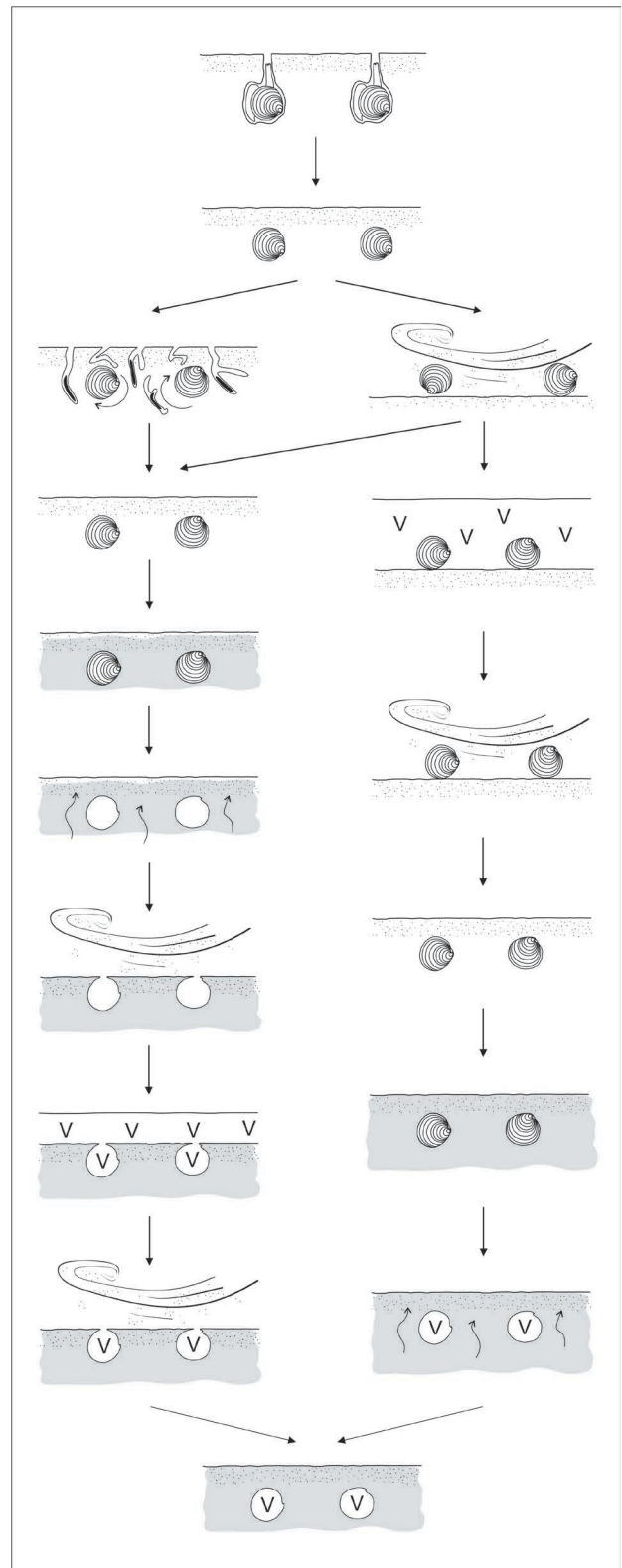


Fig. 6 - Schematic reconstruction of the taphonomic history of *Dosinia ponderosa* (Gray, 1838) in the Pisco Formation. On the left, the first scenario, on the right, the second scenario (see Discussion for the explanation of the processes). The gray halo is the consolidated sediment, the thin arrows represent the circulating diagenetic fluids, the "V" symbol indicates the volcanic ash.

the species *Dosinia ponderosa* (Gray, 1838) thanks to the fine-grained volcanoclastic material of which they are

composed. Analyses of the volcanic composition, mainly consisting of glass shards and biotite crystals, suggest that these mollusks or their molds experienced a “Pompeii-effect”, i.e., a rapid burial caused by the primary deposition of a volcanic ash layer. The succession of taphonomic processes leading to the formation of these casts remains unclear, but two main alternatives are hypothesized herein. In both cases, it was the strong explosive volcanic activity affecting the East Pisco Basin during the Late Miocene that allowed this peculiar preservation style in a shallow-water marine setting.

ACKNOWLEDGEMENTS

I would like to warmly thank A. Collareta, C. Di Celma, E. Malinverno, L. Mariani and G. Coletti for greatly contributing in realizing this work. Special thanks to M. Urbina for his assistance in the field. I also wish to thank G. Bianucci, A. Gioncada and T.J. DeVries for fruitful discussions about the stratigraphy, invertebrate paleontology and diagenesis of the Pisco Basin. Thanks to R. Varas-Malca, R. Salas-Gismondi, A. Altamirano-Sierra, and W. Aguirre for their support at the Museo de Historia Natural de la Universidad Nacional Mayor de San Marcos in Lima. I also wish to thank the Associate Editor Barbara Cavalazzi and two anonymous reviewers for their constructive suggestions. This study was supported by a grant from the Italian Ministero dell’Università e della Ricerca (PRIN Project 2022MAM9ZB) and a grant from the Università degli Studi di Milano-Bicocca (“Premio Giovani Talenti” 2020-ATESP-0121 to G.B.).

REFERENCES

- Alleman V. (1978). Estudio de los macrofósiles del Mioceno de la Costa Peruana. *Revista de la Universidad Particular Ricardo Palma*, 1: 88-116.
- Amiot R., Göhlich U.B., Lécuyer C., Muizon de C., Cappetta H., Fourel F., Héran M.-A. & Martineau F. (2008). Oxygen isotope compositions of phosphate from Middle Miocene-Early Pliocene marine vertebrates of Peru. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 264: 85-92.
- Antoine P.O., Orliac M.J., Atici G., Ulusoy I., Sen E., Çubukçu H.E., Albayrak E., Oyal N., Aydar E. & Sen S. (2012). A rhinocerotid skull cooked-to-death in a 9.2 Ma-old ignimbrite flow of Turkey. *PLoS One*, 7: e49997.
- Baales M., Jöris O., Street M., Bittmann F., Weninger B. & Wiethold J. (2002). Impact of the Late Glacial eruption of the Laacher See volcano, central Rhineland, Germany. *Quaternary Research*, 58: 273-288.
- Bamforth E.L., Narbonne G.M. & Anderson M.M. (2008). Growth and ecology of a multi-branched Ediacaran rangeomorph from the Mistaken Point assemblage, Newfoundland. *Journal of Paleontology*, 82: 763-777.
- Baqueir E. (1979). Sobre la distribución de *Megapitaria aurantiaca* (Sowerby), *M. squalida* (Sowerby) y *Dosinia ponderosa* (Gray) en relación a la granulometría del sedimento (Bivalvia: Veneridae): Nota científica. *Anales del Centro de Ciencias del Mar y Limnología Universidad Nacional Autónoma de México*, 6: 25-31.
- Bianucci G. & Collareta A. (2022). An overview of the fossil record of cetaceans from the East Pisco Basin (Peru). *Bollettino della Società Paleontologica Italiana*, 61: 19-60.
- Bianucci G., Lambert O., Urbina M., Merella M., Collareta A., Bennion R., Salas-Gismondi R., Benites-Palomino A., Post K., Muizon de C., Bosio G., Di Celma C., Malinverno E., Pierantoni P.P., Villa I.M. & Amson E. (2023). A heavyweight early whale pushes the boundaries of vertebrate morphology. *Nature*, 620: 824-829.
- Bosio G., Gioncada A., Malinverno E., Di Celma C., Villa I.M., Cataldi G., Gariboldi K., Collareta A., Urbina M. & Bianucci G. (2019). Chemical and petrographic fingerprinting of volcanic ashes as a tool for high-resolution stratigraphy of the upper Miocene Pisco Formation (Peru). *Journal of the Geological Society*, 176: 13-28.
- Bosio G., Gioncada A., Di Celma C., Villa I.M., Pichavant M., Urbina M. & Bianucci G. (2020a). Two-mica rhyolitic tephra in the East Pisco Basin (Peru): new age and dispersion constraints for the eruptions of the Eastern Cordillera of Central Andes. *Bulletin of Volcanology*, 82: 1-22.
- Bosio G., Malinverno E., Collareta A., Di Celma C., Gioncada A., Parente M., Berra F., Marx F.G., Vertino A., Urbina M. & Bianucci G. (2020b). Strontium isotope stratigraphy and the thermophilic fossil fauna from the middle Miocene of the East Pisco Basin (Peru). *Journal of South American Earth Sciences*, 97: 102399.
- Bosio G., Malinverno E., Villa I.M., Di Celma C., Gariboldi K., Gioncada A., Barberini V., Urbina M. & Bianucci G. (2020c). Tephrochronology and chronostratigraphy of the Miocene Chilcatay and Pisco Formations (East Pisco Basin, Peru). *Newsletters on Stratigraphy*, 53: 213-247.
- Bosio G., Bracchi V., Malinverno E., Collareta A., Coletti G., Gioncada A., Kočí T., Di Celma C., Bianucci G. & Basso D. (2021a). Taphonomy of a *Panopea Ménard* de la Groye, 1807 shell bed from the Pisco Formation (Miocene, Peru). *Comptes Rendus Palevol*, 20: 119-140.
- Bosio G., Collareta A., Di Celma C., Lambert O., Marx F.G., de Muizon C., Gioncada A., Gariboldi K., Malinverno E., Varas Malca R., Urbina M. & Bianucci G. (2021b). Taphonomy of marine vertebrates of the Pisco Formation (Miocene, Peru): insights into the origin of an outstanding Fossil-Lagerstätte. *PLoS One*, 16: e0254395.
- Bosio G., Gioncada A., Gariboldi K., Bonaccorsi E., Collareta A., Pasero M., Di Celma C., Malinverno E. & Bianucci G. (2021c). Mineralogical and geochemical characterization of fossil bones from a Miocene marine Konservat-Lagerstätte. *Journal of South American Earth Sciences*, 105: 102924.
- Bosio G., Bianucci G., Collareta A., Landini W., Urbina M. & Di Celma C. (2022). Ultrastructure, composition, and ⁸⁷Sr/⁸⁶Sr dating of shark teeth from lower Miocene sediments of southwestern Peru. *Journal of South American Earth Sciences*, 118: 103909.
- Briggs D.E., Siveter D.J., Siveter D.J. & Sutton M.D. (2008). Virtual fossils from 425 Million-year-old volcanic ash: a set of exceptionally preserved but difficult-to-extract fossils reveals the diverse creatures from a Silurian sea-floor community. *American Scientist*, 96: 474-481.
- Carey S. (1997). Influence of convective sedimentation on the formation of widespread tephra fall layers in the deep sea. *Geology*, 25: 839-842.
- Cherns L., Wheeley J.R. & Wright V.P. (2011). Taphonomic Bias in Shelly Faunas Through Time: Early Aragonitic Dissolution and Its Implications for the Fossil Record. In Allison P.A. & Bottjer D.J. (eds), *Taphonomy. Aims & Scope Topics in Geobiology Book Series*. Springer, Dordrecht, 32: 79-105.
- Coan E.V. & Valentich-Scott P. (2012). Bivalve seashells of tropical West America. Marine bivalve mollusks from Baja California to northern Peru. 6. 1258 pp. Santa Barbara Museum of Natural History, Santa Barbara.
- Coletti G., Bosio G., Collareta A., Malinverno E., Bracchi V., Di Celma C., Basso D., Stainbank S., Spezzaferri S., Cannings T. & Bianucci G. (2019a). Biostratigraphic, evolutionary, and paleoenvironmental significance of the southernmost lepidocyclinids of the Pacific coast of South America (East Pisco Basin, southern Peru). *Journal of South American Earth Sciences*, 96: 102372.
- Coletti G., Collareta A., Bosio G., Urbina-Schmitt M. & Buckeridge J. (2019b). *Perumegabalanus calzai* gen. et sp. nov., a new intertidal megabalanine barnacle from the early Miocene

- of Peru. *Neues Jahrbuch für Geologie und Paläontologie, Abhandlungen*, 294: 197-212.
- Collareta A., Coletti G., Bosio G., Buckeridge J., de Muizon C., DeVries T.J., Varas-Malca R., Altimirano-Sierra A., Urbina-Schmitt M. & Bianucci G. (2019). A new barnacle (Cirripedia: Neobalanoformes) from the early Miocene of Peru: Palaeoecological and palaeobiogeographical implications. *Neues Jahrbuch für Geologie und Paläontologie, Abhandlungen*, 292: 321-338.
- Collareta A., Lambert O., Marx F.G., Muizon C. de, Varas-Malca R., Landini W., Bosio G., Malinverno E., Gariboldi K., Gioncada A., Urbina M. & Bianucci G. (2021). Vertebrate palaeoecology of the Pisco Formation (Miocene, Peru): glimpses into the Ancient Humboldt Current Ecosystem. *Journal of Marine Science and Engineering*, 9: 1188.
- Collareta A., Bosio G., Varas-Malca R., Bianucci G., Merella M., Urbina M. & Di Celma C. (2023). The extinct nautiloid *Aturia* in the Middle Miocene of Pacific South America: new data from the Pisco Lagerstätte of Peru. *Neues Jahrbuch für Geologie und Paläontologie*, 308: 23-32.
- Deem J.M. (2005). *Bodies from the Ash*. 48 pp. Houghton Mifflin Harcourt, New York.
- DeVries T.J. (1986). The geology and paleontology of Tablazos in Northwest Peru. 964 pp. PhD Thesis, The Ohio State University.
- DeVries T.J. (1988). Mollusks of the Pisco Basin. *Pisco Basin Guidebook*, 8: 127-134.
- DeVries T.J. (1997). Neogene *Ficus* (Mesogastropoda: Ficidae) from the Pisco Basin (Peru). *Boletín de la Sociedad Geológica del Perú*, 86: 11-18.
- DeVries T.J. (2001). Molluscan evidence for an Oligocene-Miocene age of 'Paracas' beds in southern Peru. *Boletín de la Sociedad Geológica del Perú*, 92: 57-65.
- DeVries T.J. (2007). Cenozoic Turritellidae (Gastropoda) from southern Peru. *Journal of Paleontology*, 81: 331-351.
- DeVries T.J. (2016). Fossil Cenozoic crassatelline bivalves from Peru: New species and generic insights. *Acta Palaeontologica Polonica*, 61: 661-688.
- DeVries T.J. (2017). Eocene stratigraphy and depositional history near Puerto Caballas (East Pisco Basin, Peru). *Boletín de la Sociedad Geológica del Perú*, 112: 39-52.
- DeVries T.J. (2019). Early Paleogene brackish-water mollusks from the Caballas Formation of the East Pisco Basin (southern Peru). *Journal of Natural History*, 53: 1533-1584.
- DeVries T.J. & Frassinetti D. (2003). Range extensions and biogeographic implications of Chilean Neogene mollusks found in Peru. *Boletín del Museo Nacional de Historia Natural de Chile*, 52: 119-135.
- DeVries T.J. & Jud N.A. (2018). Lithofacies patterns and paleogeography of the Miocene Chilcatay and lower Pisco depositional sequences (East Pisco Basin, Peru). *Boletín de la Sociedad Geológica del Perú*, 8: 124-167.
- DeVries T.J., Groves L.T. & Urbina M. (2006). A new early Miocene *Muracypraea* Woodring, 1957 (Gastropoda: Cypraeidae) from the Pisco Basin of southern Peru. *Nautilus*, 120: 101-105.
- DeVries T.J., Urbina M. & Jud N.A. (2017). The Eocene-Oligocene Otuma depositional sequence (East Pisco Basin, Peru): paleogeographic and paleoceanographic implications of new data. *Boletín de la Sociedad Geológica del Perú*, 112: 14-38.
- Di Celma C., Ragaini L., Cantalamessa G. & Curzio P. (2002). Shell concentrations as tools in characterizing sedimentary dynamics at sequence-bounding unconformities: examples from the lower unit of the Canoa Formation (Late Pliocene, Ecuador). *Geobios*, 35: 72-85.
- Di Celma C., Malinverno E., Bosio G., Collareta A., Gariboldi K., Gioncada A., Mollí G., Basso D., Varas-Malca R.M., Pierantoni P.P., Villa I.M., Lambert O., Landini W., Sarti G., Cantalamessa G., Urbina M. & Bianucci G. (2017). Sequence stratigraphy and paleontology of the upper Miocene Pisco Formation along the western side of the lower Ica valley (Ica Desert, Peru). *Rivista Italiana di Paleontologia e Stratigrafia*, 123: 255-274.
- Di Celma C., Pierantoni P.P., Volatili T., Mollí G., Mazzoli S., Sarti G., Ciattoni S., Bosio G., Malinverno E., Collareta A., Gariboldi K., Gioncada A., Jablonska D., Landini W., Urbina M. & Bianucci G. (2022). Towards deciphering the Cenozoic evolution of the East Pisco Basin (southern Peru). *Journal of Maps*, 18: 1-16.
- DiMichele W.A. & Falcon-Lang H.J. (2011). Pennsylvanian 'fossil forests' in growth position (T0 assemblages): origin, taphonomic bias and palaeoecological insights. *Journal of the Geological Society*, 168: 585-605.
- Droser M.L., Gehling J.G. & Jensen S.R. (2006). Assemblage palaeoecology of the Ediacara biota: the unabridged edition? *Palaeogeography, Palaeoclimatology, Palaeoecology*, 232: 131-147.
- Dunbar R.B., Marty R.C. & Baker P.A. (1990). Cenozoic marine sedimentation in the Sechura and Pisco Basins, Peru. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 77: 235-261.
- Esperante R. & Poma O. (2015). Taphonomy and palaeopathology of two mysticete whales, upper Miocene Pisco Formation, Peru. *Spanish Journal of Palaeontology*, 30: 1-14.
- Esperante R., Brand L.R., Chadwick A.V. & Poma O. (2015). Taphonomy and paleoenvironmental conditions of deposition of fossil whales in the diatomaceous sediments of the Miocene/Pliocene Pisco Formation, southern Peru - A new fossil-lagerstätte. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 417: 337-370.
- Famoso N.A. & Pagnac D. (2011). A Comparison of the Clarendonian Equid Assemblages from the Mission Pit, South Dakota and Ashfall Fossil Beds, Nebraska. *Transactions of the Nebraska Academy of Sciences and Affiliated Societies*, 32: 98-107.
- Gariboldi K., Gioncada A., Bosio G., Malinverno E., Di Celma C., Tinelli C., Cantalamessa G., Landini W., Urbina M. & Bianucci G. (2015). The dolomite nodules enclosing fossil marine vertebrates in the East Pisco Basin, Peru: field and petrographic insights into the Lagerstätte formation. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 438: 81-95.
- Giacomelli L., Perrotta A., Scandone R. & Scarpati C. (2003). The eruption of Vesuvius of 79 AD and its impact on human environment in Pompeii. *Episodes*, 26: 235-238.
- Gioncada A., Gariboldi K., Collareta A., Di Celma C., Bosio G., Malinverno E., Lambert O., Pike J., Urbina M. & Bianucci G. (2018a). Looking for the key to preservation of fossil marine vertebrates in the Pisco Formation of Peru: New insights from a small dolphin skeleton. *Andean Geology*, 45: 379-398.
- Gioncada A., Petrini R., Bosio G., Gariboldi K., Collareta A., Malinverno E., Bonaccorsi E., Di Celma C., Pasero M., Urbina M. & Bianucci G. (2018b). Insights into the diagenetic environment of fossil marine vertebrates of the Pisco Formation (late Miocene, Peru) from mineralogical and Sr-isotope data. *Journal of South American Earth Sciences*, 81: 141-152.
- Goetcheus V.G. & Birks H.H. (2001). Full-glacial upland tundra vegetation preserved under tephra in the Beringia National Park, Seward Peninsula, Alaska. *Quaternary Science Reviews*, 20: 135-147.
- Góis-Marques C.A., De Nascimento L., Menezes de Sequeira M., Fernández-Palacios J.M. & Madeira J. (2019). The Quaternary plant fossil record from the volcanic Azores Archipelago (Portugal, North Atlantic Ocean): a review. *Historical Biology*, 31: 1267-1283.
- Grandi F., Del Valle H., Caceres I., Rodríguez-Salgado P., Oms O., Fernández-Jalvo Y., García F., Campeny G. & Gomez de Soler B. (2023). Exceptional preservation of large fossil vertebrates in a volcanic setting (Camp dels Ninots, Spain). *Historical Biology*, 35: 1234-1249.
- Gray J.E. (1838). Catalogue of the species of the genus *Cytherea*, of Lamarck, with the description of some new genera and species. *The Analyst*, 8: 302-309.

- Griggs A.J., Davies S.M., Abbott P.M., Rasmussen T.L. & Palmer A.P. (2014). Optimising the use of marine tephrochronology in the North Atlantic: a detailed investigation of the Faroe Marine Ash Zones II, III and IV. *Quaternary Science Reviews*, 106: 122-139.
- Hay R.L. (1986). Role of tephra in the preservation of fossils in Cenozoic deposits of East Africa. *Geological Society, London, Special Publications*, 25: 339-344.
- Hay R.L. & Leakey M.D. (1982). The fossil footprints of Laetoli. *Scientific American*, 246: 50-57.
- Hayward J.J. & Hayward B.W. (1995). Fossil forests preserved in volcanic ash and lava at Ihumatao and Takapuna, Auckland. *Tane*, 35: 127-142.
- Hayward J.L., Amlaner C.J. & Young K.A. (1989). Turning eggs to fossils: A natural experiment in taphonomy. *Journal of Vertebrate Paleontology*, 9: 196-200.
- Heikoop J.M., Tsujita C.J., Heikoop C.E., Risk M.J. & Dickin A.P. (1996a). Effects of volcanic ashfall recorded in ancient marine benthic communities: comparison of a nearshore and an offshore environment. *Lethaia*, 29: 125-139.
- Heikoop J.M., Tsujita C.J., Risk M.J. & Tomascik T. (1996b). Corals as proxy recorders of volcanic activity: evidence from Banda Api, Indonesia. *Palaios*, 11: 286-292.
- Hofmann H.J., O'Brien S.J. & King A.F. (2008). Ediacaran biota on Bonavista Peninsula, Newfoundland, Canada. *Journal of Paleontology*, 82: 1-36.
- Hsu J.T. (1992). Quaternary uplift of the Peruvian coast related to the subduction of the Nazca Ridge: 13.5 to 15.6 degrees south latitude. *Quaternary International*, 15: 87-97.
- Iurino D.A., Bellucci L., Schreve D. & Sardella R. (2014). Exceptional soft tissue fossilization of a Pleistocene vulture (*Gyps fulvus*): new evidence for emplacement temperatures of pyroclastic flow deposits. *Quaternary Science Reviews*, 96: 180-187.
- Jiang B., Harlow G.E., Wohletz K., Zhou Z. & Meng J. (2014). New evidence suggests pyroclastic flows are responsible for the remarkable preservation of the Jehol biota. *Nature Communications*, 5: 3151.
- Kiel S., Jakubowicz M., Altamirano A., Belka Z., Dopierska J., Urbina M. & Salas-Gismondí R. (2023). The late Cenozoic evolution of the Humboldt Current System in coastal Peru: Insights from neodymium isotopes. *Gondwana Research*, 116: 104-112.
- Kočí T., Bosio G., Collareta A., Sanfilippo R., Ekrť B., Urbina M., Malinverno E. (2021). First report on the cirratulid (Annelida, polychaeta) reefs from the Miocene Chilcatay and Pisco Formations (East Pisco Basin, Peru). *Journal of South American Earth Sciences*, 107: 103042.
- Kovács J., Németh K., Szabó P., Kocsis L., Kereszturi G., Újvári G. & Vennemann T. (2020). Volcanism and paleoenvironment of the pula maar complex: A pliocene terrestrial fossil site in Central Europe (Hungary). *Palaeogeography, Palaeoclimatology, Palaeoecology*, 537, 109398.
- Kutterolf S., Schacht U., Wehrmann H., Freundt A. & Mörz T. (2006). Onshore to offshore tephrostratigraphy and marine ash layer diagenesis. In Bunschuh J. & Alvarado G.E. (eds), *Central America: Geology, Resources and Hazards*: 395-421.
- Lambert O., Bianucci G., Post K., De Muizon C., Salas-Gismondí R., Urbina M. & Reumer J. (2010). The giant bite of a new raptorial sperm whale from the Miocene epoch of Peru. *Nature*, 466: 105-108.
- Lazo D.G. (2004). Bivalve taphonomy: testing the effect of life habits on the shell condition of the littleneck clam *Protothaca (Protothaca) staminea* (Mollusca: Bivalvia). *Palaios*, 19: 451-459.
- Lisson C.I. (1925). Algunos fósiles de Perú. *Boletín de la Sociedad Geológica del Perú*, 1: 23-30.
- Locatelli E.R. (2014). The exceptional preservation of plant fossils: a review of taphonomic pathways and biases in the fossil record. *The Paleontological Society Papers*, 20: 237-258.
- Lockley M.G. (1990). How volcanism affects the biostratigraphic record. In Lockley M.G. & Rice A. (eds), *Volcanism and fossil biotas. Geological Society of America Special Paper*, 244: 1-12.
- Lokier S.W. (2021). Marine carbonate sedimentation in volcanic settings. *Geological Society of London, Special Publications*, 520: 547-594.
- Maguire E.P. (2022). The Effect of Volcanic Ash Deposition on Marine Environments, Invertebrate Ecosystems and Fossil Preservation: Integrating Field Observations and Laboratory Experiments. 172 pp. Doctoral dissertation, Kent State University.
- Maguire E.P., Feldmann R.M., Casadio S. & Schweitzer C.E. (2016). Distal volcanic ash deposition as a cause for mass kills of marine invertebrates during the Miocene in Northern Patagonia, Argentina. *Palaios*, 31: 577-591.
- Marriott S.B., Morrissey L.B. & Hillier R.D. (2009). Trace fossil assemblages in Upper Silurian tuff beds: evidence of biodiversity in the Old Red Sandstone of southwest Wales, UK. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 274: 160-172.
- Meister P., McKenzie J.A., Vasconcelos C., Bernasconi S., Frank M., Gutjahr M. & Schrag D.P. (2007). Dolomite formation in the dynamic deep biosphere: results from the Peru Margin. *Sedimentology*, 54: 1007-1032.
- Meister P., Gutjahr M., Frank M., Bernasconi S.M., Vasconcelos C. & McKenzie J.A. (2011). Dolomite formation within the methanogenic zone induced by tectonically driven fluids in the Peru accretionary prism. *Geology*, 39: 563-566.
- Mietto P., Avanzini M. & Rolandi G. (2003). Human footprints in Pleistocene volcanic ash. *Nature*, 422: 133-133.
- Muizon de C. & Devries T.J. (1985). Geology and paleontology of late Cenozoic marine deposits in the Sacaco area (Peru). *Geologische Rundschau*, 74: 547-563.
- Narbonne G.M. (2005). The Ediacara biota: Neoproterozoic origin of animals and their ecosystems. *Annual Review of Earth and Planetary Sciences*, 33: 421-442.
- Narbonne G.M. & Gehling J.G. (2003). Life after snowball: the oldest complex Ediacaran fossils. *Geology*, 31: 27-30.
- Nelson A.E., Smellie J.L., Williams M. & Zalasiewicz J. (2008). Short Note: Late Miocene marine trace fossils from James Ross Island. *Antarctic Science*, 20: 591-592.
- Nielsen S.N. & Glodny J. (2009). Early Miocene subtropical water temperatures in the southeast Pacific. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 280: 480-488.
- Opluštil S., Pšenička J., Libertin M., Bashforth A.R., Šimůnek Z., Drábková J. & Dašková J. (2009). A Middle Pennsylvanian (Bolshevik) peat-forming forest preserved in situ in volcanic ash of the Whetstone Horizon in the Radnice Basin, Czech Republic. *Review of Palaeobotany and Palynology*, 155: 234-274.
- Orr P.J., Briggs D.E., Siveter D.J. & Siveter D.J. (2000). Three-dimensional preservation of a non-biomineralized arthropod in concretions in Silurian volcaniclastic rocks from Herefordshire, England. *Journal of the Geological Society*, 157: 173-186.
- Pfefferkorn H.W. & Wang J. (2007). Early Permian coal-forming floras preserved as compressions from the Wuda District (Inner Mongolia, China). *International Journal of Coal Geology*, 69: 90-102.
- Rivera C. (1957). Moluscos fósiles de la Formación Paracas-Ica. *Boletín de la Sociedad Geológica del Perú*, 32: 165-219.
- Rößler R. (2021). The most entirely known Permian terrestrial ecosystem on Earth-kept by explosive volcanism. *Palaeontographica Abteilung B*, 303: 1-75.
- Rößler R., Zierold T., Feng Z., Kretzschmar R., Merbitz M., Annacker V. & Schneider J.W. (2012). A snapshot of an early Permian ecosystem preserved by explosive volcanism: new results from the Chemnitz petrified forest, Germany: a snapshot of an early Permian ecosystem. *Palaios*, 27: 814-834.
- Sanfilippo R., Kočí T., Bosio G., Collareta A., Ekrť B., Malinverno E., Di Celma C., Urbina M. & Bianucci G. (2021). An investigation of vermetid reefs from the Miocene of Peru, with

- the description of a new species. *Journal of South American Earth Sciences*, 108: 103233.
- Scott A.C. (1990). Preservation, evolution and extinction of plants in Lower Carboniferous volcanic sequences in Scotland. In Lockley M.G. & Rice A. (eds), *Volcanism and fossil biotas. Geological Society of America Special Paper*, 244: 25-38.
- Seilacher A. (1992). Vendobionta and Psammocorallia: lost constructions of Precambrian evolution. *Journal of the Geological Society*, 149: 607-613.
- Thornburg T.M. & Kulm L.D. (1981). Sedimentary basins of the Peru continental margin: Structure, stratigraphy, and Cenozoic tectonics from 6 S to 16 S latitude. *Nazca plate: Crustal formation and Andean convergence*, 154: 393-422.
- Villaseca A., Champi N. & Pickling J. (2020). Fossil register of the Paleontological Didactic Museum of Ocucaje, Ica, Perú. *Biotempo*, 17: 237-243.
- Wagner R.H. (1989). A late Stephanian forest swamp with *Sporangiostrobus* fossilized by volcanic ash fall in the Puertollano Basin, central Spain. *International Journal of Coal Geology*, 12: 523-552.
- White J.D.L. & Houghton B.F. (2006). Primary volcaniclastic rocks. *Geology*, 34: 677-680.
- Wick S.L. (2023). A new assemblage of fossil leaves and wood from The Aguja Formation (Upper Campanian) of West Texas: An interfluvial community in a volcanic landscape. *Cretaceous Research*, 142: 105366.
- Xu F., You X., Li Q. & Liu Y. (2019). Can Primary Ferroan Dolomite and Ankerite Be Precipitated? Its Implications for Formation of Submarine Methane-Derived Authigenic Carbonate (MDAC) Chimney. *Minerals*, 9: 413.
- Manuscript submitted 26 February 2023
Revised manuscript accepted 24 April 2023
Published online 31 October 2023
Editor Barbara Cavalazzi