



ELSEVIER

Contents lists available at SciVerse ScienceDirect

Comptes Rendus Palevol

www.sciencedirect.com



General palaeontology, systematics and evolution (Palaeoichnology)

Determination of a Late Miocene rocky palaeoshore by bioerosion trace fossils from the Bozcaada Island, Çanakkale, Turkey

Détermination d'un paléorivage rocheux du Miocène supérieur par des traces fossiles de bio-érosion dans l'île de Bozcaada, Çanakkale, Turquie

Huriye Demircan

Department of Geological Research, General Directorate of Mineral Research and Exploration (MTA), 06520, Ankara, Turkey

ARTICLE INFO

Article history:

Received 26 May 2011

Accepted after revision 18 November 2011

Available online 24 May 2012

Presented by Philippe Taquet

Keywords:

Bioerosion
Rocky-shore
Trace fossils
Late Miocene
Bozcaada
Turkey

Mots clés :

Bio-érosion
Rivage rocheux
Traces fossiles
Miocène supérieur
Bozcaada
Turquie

ABSTRACT

Bioerosion is a common process in hard substrates. This study introduces an example from the rocky palaeoshore cropping out at a sea cliff on the Bozcaada Island. It includes bioerosion trace fossils preserved in limestone boulders of the shallow marine and lacustrine Alcitepe Formation of Late Miocene age. The ichnotaxa include borings produced by duraphagous drillers (*Oichnus* isp.), phonorids (cf. *Conchotrema* isp.), clionid sponges (*Entobia* cf. *goniodes*, *Entobia geometrica*, *Entobia laquea*, *Entobia ovula*, *E. cf. solaris*, *Entobia* isp.), endolithic bivalves (*Gastrochaenolites torpedo*, *Gastrochaenolites lapidicus*, *Gastrochaenolites* isp., *Phrixichnus* isp.), polychaete annelids (*Maeandropolydora* isp., *Maeandropolydora sulcans*, *Maeandropolydora decipiens*, *Caulostrepis taeniola*, *Caulostrepis* isp.), echinoids (cf. *Circolites* isp.) and spinulid worms (cf. *Trypanites* isp.). Barnacles are also common as encrusters. The borings can be ascribed to the *Gastrochaenolites-Entobia* assemblage, which is typical of Neogene rocky-shores. They belong to the *Entobia* ichnofacies indicating various conditions of light, energy, and depth. Therefore they can reveal environmental changes and play an important role in forming palaeo-rocky shores and wave-cut platforms during marine transgressive events.

© 2012 Académie des sciences. Published by Elsevier Masson SAS. All rights reserved.

R É S U M É

La bio-érosion est un processus commun dans les substrats durs. L'étude présentée ici introduit un exemple de paléorivage rocheux affleurant au niveau d'une falaise marine sur l'île de Bozcaada. Il inclut des traces fossiles de bio-érosion conservées dans des blocs calcaires de la Formation Alcitepe lacustre et marine peu profonde, d'âge Miocène supérieur. Les ichnotaxa comportent des creusements provoqués par des organismes foreurs duriphages (*Oichnus* isp.), phonoriés (cf. *Conchotrema* isp.), éponges clionidées (*Entobia* cf. *goniodes*, *Entobia geometrica*, *Entobia laquea*, *Entobia ovula*, *E. cf. solaris*, *Entobia* isp.), bivalves endolithiques (*Gastrochaenolites torpedo*, *Gastrochaenolites lapidicus*, *Gastrochaenolites* isp., *Phrixichnus* isp.), annélides polychètes (*Maeandropolydora* isp., *Maeandropolydora sulcans*, *Maeandropolydora decipiens*, *Caulostrepis taeniola*, *Caulostrepis* isp.), échinoïdes (cf. *Circolites* isp.) et vers spinulidés (cf. *Trypanites* isp.). Les barnacles sont aussi des organismes encroûtants. Les creusements peuvent être attribués à l'assemblage *Gastrochaenolites-Entobia*, typique des rivages rocheux néogènes. Ils appartiennent à l'ichnofaciès *Entobia* indiquant

E-mail address: asmin68@yahoo.com.tr

des conditions variées de lumière, d'énergie et de profondeur. C'est pourquoi, ils peuvent révéler des changements environnementaux et jouer un rôle important dans la formation de paléorivages rocheux et de plates-formes d'érosion marine au cours des événements marins transgressifs.

© 2012 Académie des sciences. Publié par Elsevier Masson SAS. Tous droits réservés.

1. Introduction

Bioerosion, designed by Neumann (1966) as an abbreviation of 'biological erosion', identifies every form of biological erosion of hard substrates, including lithic and woody substrates (Bromley, 1992). Organisms can erode hard substrates by mechanical or chemical means, resulting in the formation of scrape traces, borings and other biogenic structures. Encrustation is the behaviour by which organisms attach to hard substrates. Encrusting and boring organisms represent a community that changes with the substrate that sustains them (Hohman, 1993).

Rocky-shores, as hard surfaces exposed in shallow marine environments associated with null or low sedimentation, offer excellent conditions for colonization by marine bioeroding organisms (Cachão et al., 2008). These environments represent a unique marine sedimentary environment where wave-cut marine erosion platforms are formed. Therefore, the identification of rocky-shores in the geological record is very important, because they represent major flooding surfaces and provide us with crucial information about palaeoshorelines and sea level changes. Bioerosional structures play a major role in their recognition (Ghibaudo et al., 1996; Jia-Yu and Johnson, 1996; Martinell and Domènech, 1995; Santos et al., 2008).

Bioerosional evidence is also excellent for the study of the evaluation of marine communities in these types of environments (Johnson, 1992) since their remains are often preserved in situ, connected to the hard surface itself (Brett, 1988; Johnson, 1992; Taylor and Wilson, 2003). The case of ancient marine hard substrate communities has been described from the Lower Cambrian (e.g. Kobluk, 1981a, b; Kobluk and James, 1979; Kobluk et al., 1978) to the Neogene (e.g. Bromley and Asgaard, 1993a, b; Bromley and D'Alessandro, 1987; Domènech et al., 2001; Gibert et al., 1998; Martinell and Domènech, 1995; Radwański, 1970; Watkins, 1990).

Although bioerosional structures are very important in revealing the effects of biological factors on coastal morphology, there are very few publications on bioerosion in Turkey. The first paper related to bioerosion in Turkey, 'Neogene borings from rocky-shore', is Uchman et al. (2002), who explained the relative sea level changes recorded in borings from a Miocene rocky-shore of the Mut Basin, southern Turkey. The present study uses bioerosional structures to contribute to the interpretation of palaeoenvironmental conditions in the rocky-shore of the Late Miocene depositional sequence of the Alcitepe Formation. It firstly focuses on the identification and the description of bioerosion structures with their tracemakers and then analyses the borings in terms of their substrate nature.

2. Geological setting

The studied rocks of the Alcitepe Formation are located along the southwestern coast of Bozcaada Island (Fig. 1A–C), on the southern edge of the Thracian Basin (e.g. Göncüoğlu, 2010). Previous palaeontological and sedimentological studies on the Neogene deposits of the island are quite limited (e.g. Çağatay et al., 2006; Druit, 1961; Kesgin and Varol, 2003; Önem, 1974; Sakiñç et al., 1999; Siyako et al., 1989; Temel and Çiftçi, 2002). These Neogene sediments are composed of various coeval lithofacies with lateral and vertical transitions (Dermitzakakis and Papanikolaou, 1981; Sakiñç and Yaltrak, 1997; Sakiñç et al., 1999; Yaltrak, 1996; Yaltrak and Alpar, 2002; Yaltrak et al., 1998, 2000).

Regarding previous studies, the lowermost unit in the generalized stratigraphical section (Fig. 2) is represented by the Eocene–Oligocene Ayvacik Volcanics, mainly comprising andesitic lavas and pyroclastics. Pillow lavas at some localities characterize quiescent submarine volcanic activity. The Ayvacik Volcanics are disconformably overlain by the Gazhanedere Formation, which is characterized by mudstone and conglomeratic sandstone (Middle Miocene–Lower Pliocene), representing a fluvial and lagoonal environment. On top of the Gazhanedere Formation, with lateral and vertical transitions, rests the Kirazli Formation. This consists of massive and semi-consolidated sandstone representing beach and back-beach lithofacies of the Late Miocene (Sakiñç et al., 1999). Both formations are overlain by the Alcitepe Formation (Late Miocene) with lateral and vertical transitions. This formation consists of bioclastic and oolitic limestone with basal clastic rocks, and contains molluscan and ostracod faunas in the western outcrops. These faunas indicate deposition in shallow, brackish to fresh water environments. On the other hand, previous studies (Çağatay et al., 2006) on faunal and palaeomagnetic analyses of the Alcitepe Formation in other parts of the Aegean Sea show that the formation represents chron C3r (6.04–5.24 Ma). The ostracod analysis also indicates that during the deposition of the Alcitepe Formation salinity increased from brackish in the lower part to more saline conditions in the upper part (Çağatay et al., 2006).

In the study area, a non-depositional and denudational period continued until the Pleistocene. During the transgression in the Middle–Late Pleistocene (Yaltrak et al., 2002), the valleys and broad coastal plains were affected by warm sea conditions. The terrace deposits formed during this transgression include conglomerates and sandstones, which are rich in *Ostrea edulis* shells (Sakiñç et al., 1999; Yaltrak, 1996; Yaltrak et al., 2002). In general, they overlie the Miocene rocks with an angular unconformity in this area.

The borings described in this article occur within the Alcitepe Formation.

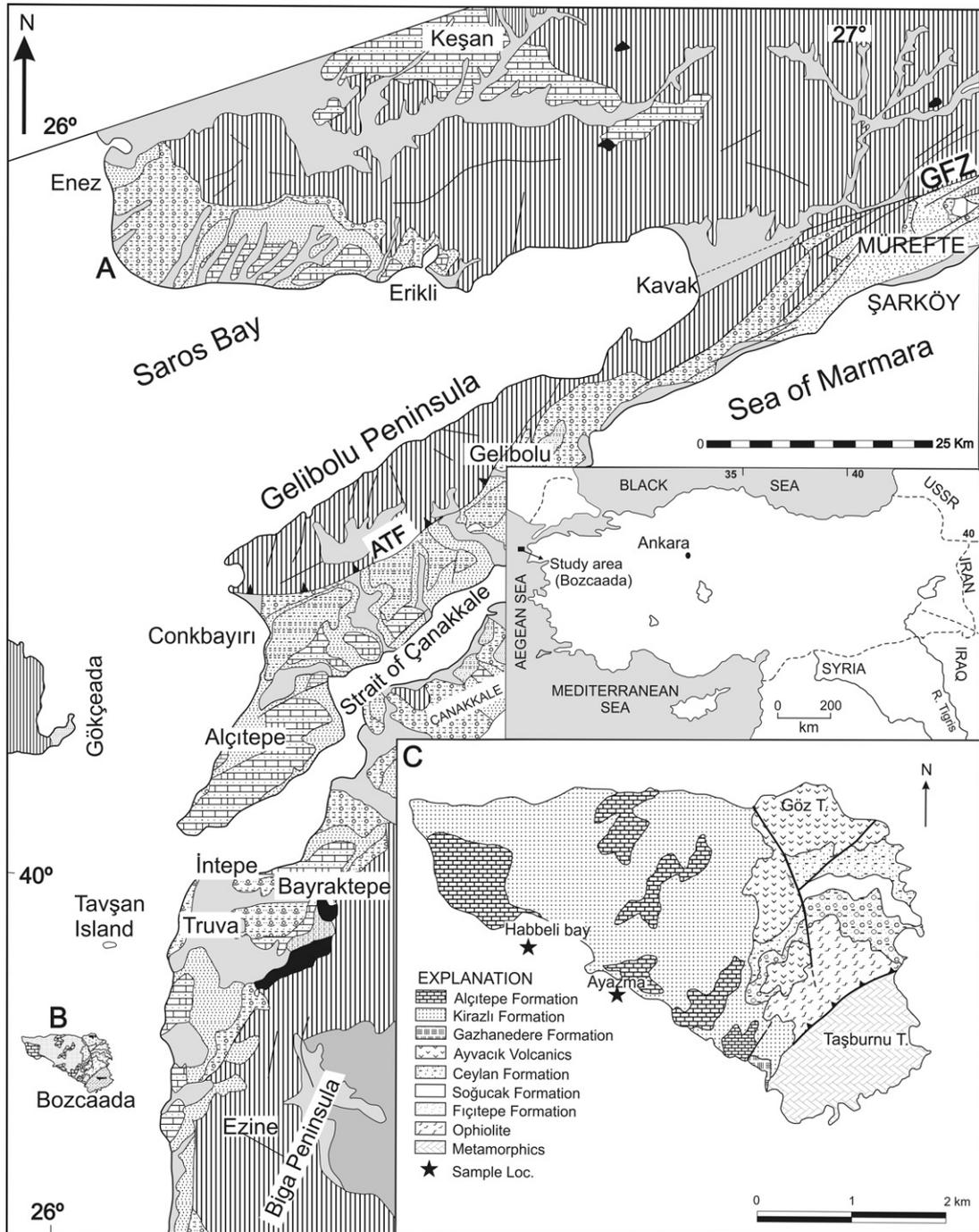


Fig. 1. A, B. Geological map of the northeastern Aegean region (modified from Sakinç and Yaltrak, 2005). C. Close up view of the study area (modified from Temel and Çiftçi, 2002).

Fig. 1. A, B. Carte géologique de la région égéenne nord-orientale (modifié selon Sakinç et Yaltrak, 2005). C. Encart montrant la zone d'étude (modifié selon Temel et Çiftçi, 2002).

3. Description of the locality

The Alçıtepe Formation is quite extensive in Bozcaada Island and shows up as elevations and hills due to its physical features (e.g. erosion-resistance). The formation shows different characteristics from place to place. For instance,

along the northern coast of the Gulf of Saros, the formation consists of *Maetra*-bearing bioclastic and cross-bedded limestones, interbedded with *Cardium*-bearing sandstones and *Ostrea* banks. These sediments indicate fluctuations in salinity from brackish water to normal marine conditions. In the Gelibolu and Biga peninsulas the formation consists

AGE	LITHOLOGY AND FORMATION	EXPLANATION
QUATERNARY		Mud, clay with marine shells, laterally passing to alluvial deposits
LATE PLOIOCENE	3.4-3.7 My	Erosional period (starting of the North Anatolian Fault)
MIDDLE MIOCENE LOWER PLOIOCENE	ALÇITEPE FM KIRAZLI FM GAZHANEDERE FM	Limestone-mud intercalations bivalve and gastropoda (lacustrine and shallow marine environments) Massive sandstone bivalve and gastropoda (beach and back-beach environments)
Eocene OLIGOCENE	AYVACIK	Sandstone-shale Mainly andezidic lava and pyroclastics

(not to scale)

Fig. 2. Generalized measured section of the study area (modified from Sakiç and Yaltrak, 2005).

Fig. 2. Section générale de la zone étudiée (modifié selon Sakiç et Yaltrak, 2005).

of mudstone and marl in the lower part, and bioclastic and oolitic limestones with marl, mudstone, and sandstone intercalations in the upper part (Çağatay et al., 2006).

The best exposure of the Alcitepe Formation in the studied area is located along Habbeli Bay in the southwestern part of the island (Fig. 1C), between N 39° 49' 107; E 025° 59' 026 and N 39° 49' 2877; E 025° 59' 369, along the cliff of Ayazma beach in Habbeli Bay, and southeast of the few inland localities which are represented by carbonate and siliciclastic units (Figs. 3 and 4).

At Habbeli Bay, the Alcitepe Formation is approximately 30 metres thick and rests on the Kirazli Formation. The formation starts with *Mastra*-bearing, beige-grey coloured, medium-bedded, pebbly to coarse-grained sandy limestone, and continues with conglomerates alternating with sandstones, succeeded by marls (18 cm thickness) in the study area. The marls of the formation contain a brackish water fauna with *Mastra* shells (Fig. 5).

Towards the top of the section, *Ostrea*- and *Pecten*-bearing calcareous marine sandstones occur. Directly on top of this level is a 70-cm thick coquin composed of bivalve and gastropods shells, which can be traced as a marker. The bivalves are almost completely dissolved with only their moulds remaining. The unit continues as whitish grey coloured, thin-medium layered limestone, sandy limestone and mudstone towards the top.

Oolitic limestone can be observed in some localities, as can large-scaled tabular cross stratification (Fig. 6) and lightly cemented spherical sand clumps. The presence of large-scaled cross stratification and widespread ooids



Fig. 3. Wave-cut platform produced by wave erosion in the Alcitepe Formation (Habbeli Bay).

Fig. 3. Plate-forme d'érosion marine produite par l'érosion des vagues dans la Formation Alcitepe (Baie d'Habbeli).



Fig. 4. General view of the Alcitepe Formation (Habbeli Bay).

Fig. 4. Vue générale de la formation Alcitepe (Baie d'Habbeli).



Fig. 5. *Maetra* shells in the grey coloured marls at Alcitepe Formation (Bozcaada Island, Turkey).

Fig. 5. Coquilles de *Maetra* dans les marnes grises de la Formation Alcitepe (île de Bozcaada, Turquie).



Fig. 6. Shallow marine-shore face cross lamination, Alcitepe Formation.

Fig. 6. Stratification entrecroisée sur une face de rivage marin peu profond, Formation Alcitepe.

indicates high-energy levels with agitated shallow marine or lacustrine deposition progressing towards the open sea.

Towards the top of the section, *Ostrea*- and *Pecten*-bearing calcareous marine sandstone occurs. The uppermost part of the formation is made up of an alternation of *Maetra*-bearing limestones and sandstones containing *Ostrea*, followed by another alternation of *Maetra*- and *Ostrea*-bearing limestones. The presence of *Ostrea* the upper part of the Alcitepe Formation indicates normal marine depositional conditions.

The section from the middle to top of the succession is rich in fossils, represented by bivalves (Fig. 7: 1–9; 12), some echinoids (Fig. 7: 10, 11) and gastropods (Fig. 7: 13, 14). Bivalves produce *Gastrochaenolites torpedo* (Fig. 8), and *Gastrochaenolites lapidicus* is produced by some bivalves of the genera *Lithophaga*. They all are observed on a heavily karstified and brecciated Miocene carbonate hard-substrate bioeroded surface, on limestone boulders and cobbles.

In the study area, the most conspicuous trace fossil is *Gastrochaenolites*, which was produced by bivalve bioerosion preserved in epirelief and hyporelief (*G. torpedo* and *G. lapidicus*), and on the boulder surfaces (Fig. 9a, b). *Gastrochaenolites lapidicus* casts (Fig. 10) and clavate borings *Phryxichnus*-like *Gastrochaenolites* have also been recognized. These borings are perpendicular to the surface, are truncated, and some include the trace-producing organisms (Fig. 11).

Entobia is the second dominant trace fossil that is mostly observed from penetrating outer and inner parts of shells and boulders. In some cases, it is mostly seen together with *Gastrochaenolites*. The polychaete annelid borings *Caulostrepsis* and *Maeandropolydora* are observed occasionally. *Trypanites* was identified in the substrate from the walls of *G. torpedo*. Also, some skeletons of encrusting worms were frequently observed.

4. Relationship between substrate and borings

The bioerosional structures in the study area occur together with fossil assemblages including bivalves, gastropods, echinoids and also remains of encrusting organisms. Their traces are widely distributed within carbonate boulders, and shells of ostreids, and were formed by the activities of macrobioeroders. The borings are preserved in concave hyporelief and epirelief morphology, and display intense biogenic activity with abundant skeletal remains of endo and epilithic organisms. Most of the borings are common on the upper parts or the lateral surfaces of the boulders. They are perpendicular to the surface and are proximally truncated. The outer parts of the borings have been destroyed by the marine erosion. Their ethological categories, tracemaker interpretations and the relationship between the substrates and borings in the study area are summarized in Table 1.

5. Taxonomic description

The Late Miocene rocky-shore of the studied area is intensively affected by bioerosion. Bioerosional structures are widely distributed on the tops of the limestone boulders

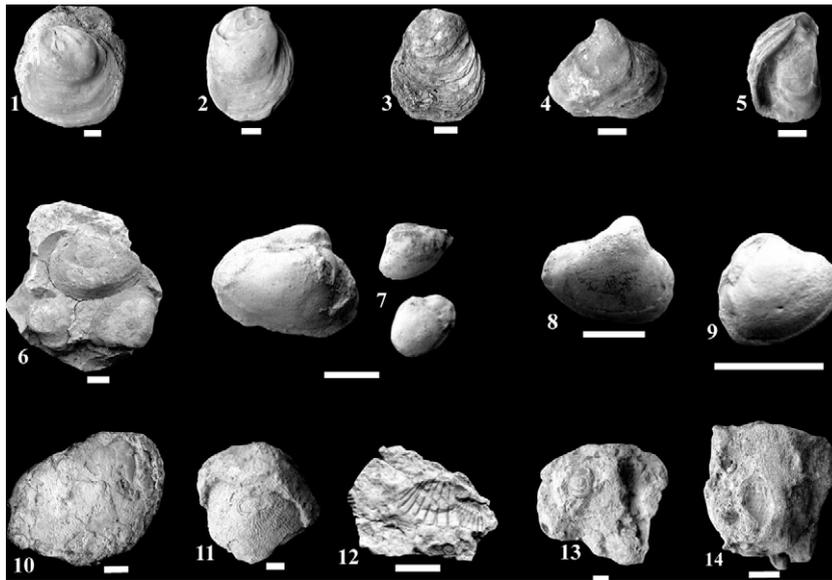


Fig. 7. Well-preserved marine macrofossils at Alcitepe Formation (Scale: 1 cm).

Fig. 7. Macrofossiles marins bien conservés de la Formation Alcitepe (échelle = 1 cm).

and on shells. The most common bioerosional traces belong to macrobioeroders, e.g., phoronids (*Conchotrema*), polychaete worms (*Caulostrepsis*, *Maeandropolydora*), clionid sponges (*Entobia*), and boring bivalves (*Gastrochaenolites*).

Nine ichnogenera have been recognized. Taxonomic description is prepared according to their ethology and morphology distinguished by Bromley (1992), Ekdale et al. (1984), Martinell (1989), and recently, Taylor and Wilson (2002). All material figured in this article is housed in the Natural History Museum, General Directorate of Mineral

Research and Exploration, MTA Ankara, Turkey (B1. 09).

Conchotrema Teichert, 1945

cf. *Conchotrema* isp. (Plate 1, Fig. A)

Description: A system of straight to slightly curved branched grooves. The grooves are 1.5–2, 0.1–0.25 mm in wide. The surfaces of the boulders were cut by their dense scrapings and they cross each other.

Remarks: Generally, these are subsurface borings, which have been exposed at the surface by erosion.

Table 1

Relationship between substrate and borings, identified at the Habbeli Bay/Ayazma beach (LB: Limestone Boulders; S: Shell).

Tableau 1

Relation entre substrat et creusements, déterminée sur l'exemple de la baie d'Habbeli/plage d'Ayazma (LB : blocs calcaires ; S : coquille).

Ichnotaxon	Tracemaker	Substrate	Ethological category
cf. <i>Conchotrema</i> isp.	Endolithic phoronid	LB	Domichnia
<i>Caulostrepsis</i> isp.	Polychaete	LB/S	Domichnia
<i>C. taeniola</i>	Polychaete	LB/S	Domichnia
<i>Circolites</i> isp.	Echinoid	LB	Domichnia
<i>Maeandropolydora</i> isp.	Polychaete	LB/S	Domichnia
<i>M. sulcans</i>	Polychaete	LB/S	Domichnia
<i>M. decipiens</i>	Polychaete	LB/S	Domichnia
<i>Entobia</i> isp.	Clionid sponge	LB/S	Domichnia
<i>E. geometrica</i>	Clionid sponge	LB	Domichnia
<i>E. cf. gonioides</i>	<i>Cliona viridis</i>	LB	Domichnia
<i>E. laquea</i>	<i>Cliona</i>	LB	Domichnia
<i>E. ovula</i>	<i>Cliona</i>	LB	Domichnia
<i>E. cf. solaris</i>	<i>Cliona</i>	LB	Domichnia
<i>Gastrochaenolites</i> isp.	Bivalve	LB/S	Domichnia
<i>G. lapidicus</i>	<i>Lithophaga</i>	LB/S	Domichnia
<i>G. torpedo</i>	<i>Lithophaga</i> & <i>Gastrochaena</i>	LB/S	Domichnia
<i>Phrixichnus</i> isp.	–	LB	Domichnia
<i>Oichnus</i> isp.	Carnivorous gastropods & Cephalopods	S	Praedichnia
<i>Trypanites</i> cf.	<i>Spinuliids</i>		Domichnia

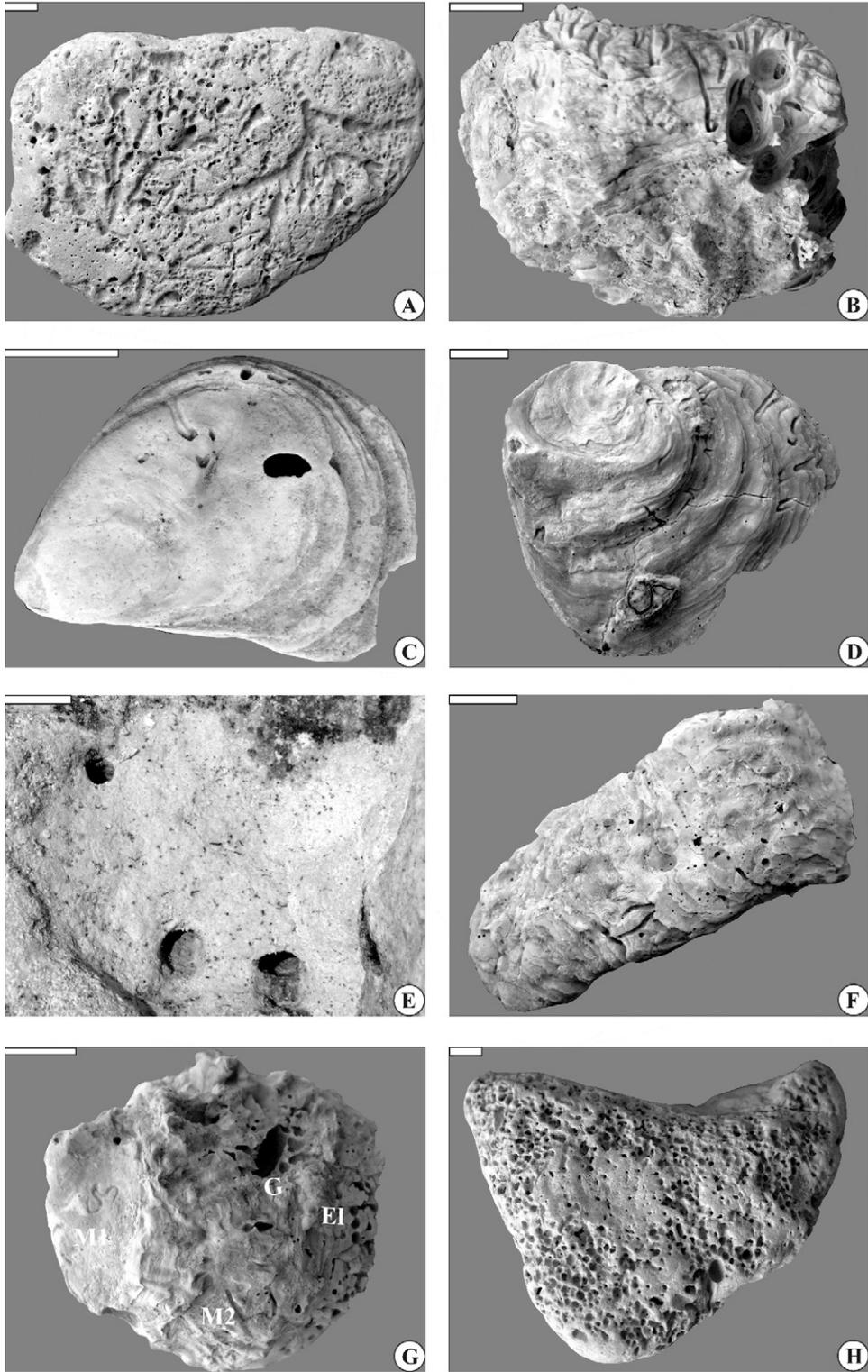


Plate 1.



Fig. 8. Plan view of *Gastrochaenolites torpedo*.
Fig. 8. Vue de *Gastrochaenolites torpedo*.

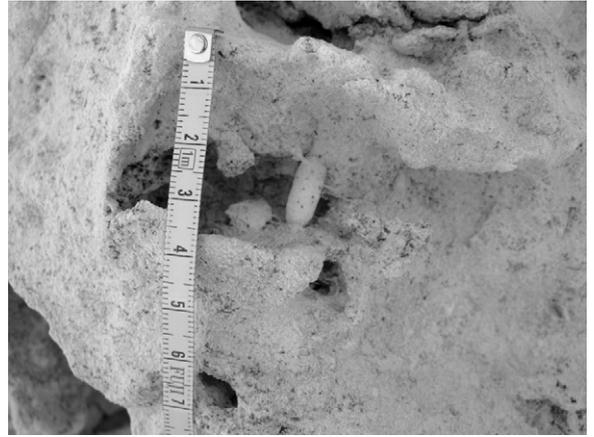


Fig. 10. Casts of *Gastrochaenolites lapidicus*.
Fig. 10. Moulages de *Gastrochaenolites lapidicus*.

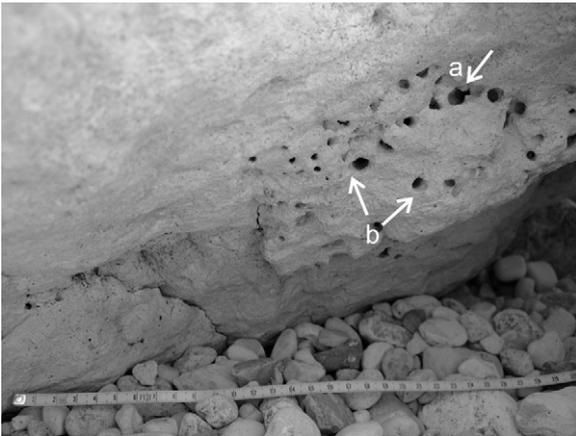


Fig. 9. View of *Gastrochaenolites torpedo* (a), *Gastrochaenolites lapidicus* (b).
Fig. 9. Vue de *Gastrochaenolites torpedo* (a) et de *Gastrochaenolites lapidicus* (b).

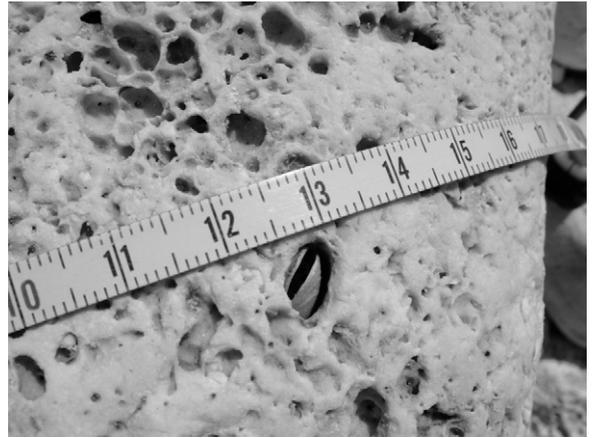


Fig. 11. *Gastrochaenolites* with its producer inside.
Fig. 11. *Gastrochaenolites* avec l'organisme à l'intérieur du trou.

Voigt (1975) suggested that *Conchotrema* is produced by endolithic phoronids.

***Caulostrepsis* Clarke, 1908**

***Caulostrepsis* isp. (Plate 1, Fig. B)**

Description: *Caulostrepsis* is a pouch-shaped boring resembling a very tight U-gallery with limbs. The limbs merge towards the aperture. The aperture is 1 mm in diameter.

Remarks: Bromley and D'Alessandro (1983) revised the systematics of this ichnotaxon and recognized several ichnospecies. It is produced by polychaetes (Radwański, 1969).

***Caulostrepsis taeniola* Clarke, 1908**

(Plate 1, Fig. C, D)

Description: This is a smooth, elongate and narrow U-shaped gallery with distinct limbs and an interconnecting

vane. The trace fossils range from 5 mm long, 0.5 mm wide (Fig. C), to 7 mm long, 0.8 mm in wide (Fig. D).

Remarks: This trace is seen on the exterior surfaces of *Ostrea* shells. *Caulostrepsis* is mainly considered to be produced by the polychaetes of the genus *Polydora* (Radwański, 1969).

Caulostrepsis occurs in shallow water environments, e.g., *Caulostrepsis cretacea* Voigt, 1971 was found between 7 and 15 m water depth (Wissihak et al., 2005).

***Circolites* Mikuláš, 1992**

cf. *Circolites* isp. (Plate 1, Fig. E)

Description: *Circolites* is a subcircular, bowl-shaped boring 3 to 4.5 cm in diameter.

Remarks: This boring is interpreted as a dwelling structure of regular echinoids (Warme, 1975), and is always restricted to very shallow waters.

***Maeandropolydora* Voigt, 1965**

***Maeandropolydora* isp. (Plate 1, Fig. F)**

Description: *Maeandropolydora* occurs as long, shallow and sinuous borings on boulders and shells that extend like 'U'-shaped pouches in the study area. It is 0.2 mm in diameter.

Remarks: This boring is produced by polychaetes of various families (Bromley and D'Alessandro, 1983).

***Maeandropolydora sulcans* Voigt, 1965**

(Plate 1, Fig. G)

Description: *M. sulcans* is a sinuous, cylindrical boring found in *Ostrea* shells in the study area small in diameter (0.5 mm) and with a narrow vane.

Remarks: It is usually attributed to annelid worms (Bromley and D'Alessandro, 1983). *M. sulcans* is commonly preserved in semirelief on rock or shell surfaces as a result of the erosion of very shallow borings.

***Maeandropolydora decipiens* Voigt, 1965**

(Plate 1, Fig. G)

Description: A single specimen is preserved on a small fragment of an oyster shell and comprises a sinuous, looped and continuous, smooth, unbranched, internally structureless gallery that parallels but clearly penetrates the shell surface. The looped limbs do not touch or intersect and the intervening oyster shell material is unaffected. Diameter of the gallery, relatively constant at both levels, is approximately 0.5 mm.

Remarks: Though incomplete, the specimen can clearly be assigned to the ichnotaxon *Maeandropolydora*, discussed in detail by Bromley and D'Alessandro (1983). *Maeandropolydora* is interpreted as the domicnion of suspension-feeding polychaete annelids (Bromley, 1994).

***Entobia* Bronn, 1837**

***Entobia* isp.** (Plate 1, Fig. H)

Description: This boring shows camerate morphology and consists of networks of chambers in planar arrangement. The shapes of chambers are from subrounded to subrectangular. The chambers are 3 mm wide.

Remarks: Entobian borings are produced by endolithic sponges. Such borings can be made by several species of *Cliona* (Bromley and D'Alessandro, 1989).

***Entobia* cf. *goniodes* Bromley and Asgaard, 1993a**

(Plate 2, Figs. A, B)

Description: This boring is observed as a system of small and camerate to nodular chambers. The chambers are 1–1.5 mm in diameter.

Remarks: *E. goniodes* is produced today by *Cliona viridis* and rarely *Cliona schmidtii* in the photic zone of the Mediterranean Sea. *C. viridis* is found at the 20 m water depth (Bromley and Asgaard, 1993a).

***Entobia laquea* Bromley and D'Alessandro, 1984**

(Plate 2, Figs. C, D)

Description: This bioerosional structure is composed of a system of tunnels and chambers in well-developed growth stages A and C sensu Bromley and D'Alessandro

(1984). It is seen here as growth stage C and is represented by irregular, oval, elongate to subangular chambers, 1.5–2 mm in diameter.

Remarks: This ichnospecies covers the surface of boulders. *Entobia laquea* is produced by sponges of the genus *Cliona* in the photic zone of the Mediterranean Sea at the present day (Bromley and Asgaard, 1993a).

***Entobia ovula* Bromley and D'Alessandro, 1984**

(Plate 2, Fig. E)

Description: This trace fossil is preserved in the A to C growth stages exposed on the surface of the boulders. The structures of stage A occur as a narrow tunnel system, with tunnels less than 1 mm in diameter. The structures of stage B are composed of curved rows of elongate chambers that are 2 mm in wide. Stage C is developed as oval, closely spaced chambers, which are 3 mm wide.

Remarks: The ichnogenus *Entobia* is produced mostly by sponges of the genus *Cliona*. Its taxonomy was discussed by Bromley and D'Alessandro (1984). *Entobia ovula* is produced in the Mediterranean today by *C. schmidtii*, *Cliona vermifera* or *Cliona vastifica* (Bromley and Asgaard, 1993a).

***Entobia* cf. *solaris* Mikuláš, 1992**

(Plate 2, Fig. F)

Description: This boring is preserved as an irregularly hemispherical depression. It also has straight radiating tunnels. The tunnels are about 1 mm in width.

Remarks: *Entobia solaris* has been described from the Late Cretaceous of the Czech Republic (Mikuláš, 1992). These borings are produced by sponges.

***Entobia geometrica* Bromley and D'Alessandro, 1984**

(Plate 2, Figs. G, H)

Description: This is a camerate ichnospecies. Apertures are of two distinct size groups. The larger apertures are circular and 4–5 mm in diameter, the smaller are between 1 mm and 2 mm in diameter.

Remarks: *E. geometrica* differs from *E. cretacea* in several respects (Bromley and D'Alessandro, 1984). The apertures of *E. geometrica* are larger and the chambers to a greater extent leave thinner dividing walls than *E. cretacea* in which the chambers are normally connected by single intercameral canals.

***Gastrochaenolites* Leymerie, 1842**

***Gastrochaenolites* isp.** (Plate 3, Fig. A)

Description: *Gastrochaenolites* is a flask-shaped boring with a narrow aperture and larger, round, elongate, straight, ovoid chambers. It is preserved as large, elongate, straight openings, steeply inclined to the host substrate. Lengths are up to 49 mm and a single observed opening is 5 mm in wide.

Remarks: This ichnogenus is produced by boring bivalves (Kelly and Bromley, 1984) such as Pholadidae, Gastrochaenidae, and Mytilidae (Fisher, 1990a, b; Warme, 1975).

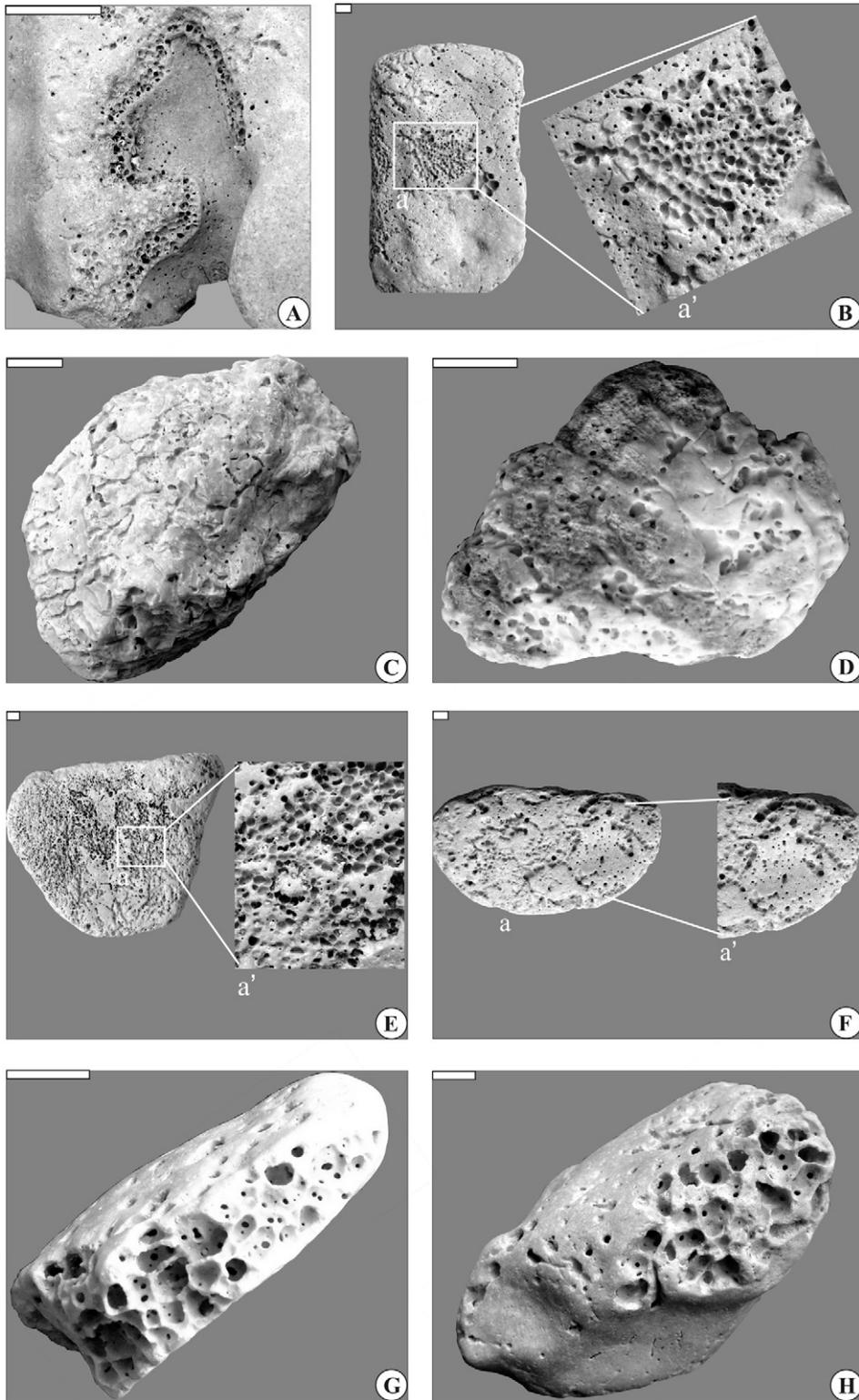


Plate 2. Field photographs of inferred clionid sponge borings from the Alcitepe limestone boulders and shells (All scales: 1 cm). **A.** *Entobia cf. goniodes*. **B.** *Entobia goniodes*. **C.** *Entobia laquea*. **D.** *Entobia laquea*. **E.** *Entobia ovula*. **F.** *Entobia cf. solaris*. **G.** *Entobia geometrica*. **H.** *Entobia geometrica*.
Planche 2. Photos sur le terrain de creusements supposés d'éponge clionidée, dans des blocs et coquilles calcaires de la Formation Alcitepe (toutes échelles = 1 cm). **A.** *Entobia cf. goniodes*. **B.** *Entobia goniodes*. **C.** *Entobia laquea*. **D.** *Entobia laquea*. **E.** *Entobia ovula*. **F.** *Entobia cf. solaris*. **G.** *Entobia geometrica*. **H.** *Entobia geometrica*.

***Gastrochaenolites lapidicus* Kelly and Bromley, 1984**
(Plate 3, Fig. B)

Description: *G. lapidicus* displays a rounded bottom. It is a smooth ovate chamber with an apertural neck and is circular throughout cross-section. The neck is also circular in cross-section or elliptical. The boring nearly 6 mm in diameter.

Remarks: This type of boring is produced by some bivalves of the genus *Lithophaga* (Kelly and Bromley, 1984).

***Gastrochaenolites torpedo* Kelly and Bromley, 1984**
(Plate 3, Figs. C, D)

Description: This is a smooth, strongly elongate chamber, 49 mm long, up to 17 mm in depth and 8–16 mm in diameter (Fig. C).

Remarks: *G. torpedo* may show a calcite lining (Jones and Pemberton, 1988) which is not observed in the studied material. This type of borings is produced by some bivalves of the genus *Lithophaga* and *Gastrochaena* (Kelly and Bromley, 1984), and in the Mediterranean region by *Lithophaga lithophaga* (Linnaeus). *G. torpedo* has been reported from Miocene rocky-shores of many regions in Europe and neighbouring areas (Radwański, 1969).

***Phrixichnus* Bromley and Asgaard, 1993a**

***Phrixichnus* isp.** (Plate 3, Fig. E)

Description: This is a smooth, strongly elongated clavate chamber. The chamber is 24 mm long and 13 mm in diameter. Its clavate shape is clearly seen.

Remarks: A clavate, *Gastrochaenolites*-like boring but exhibiting a very particular ornamentation on the walls consisting of arcuate or concentric grooves in two gently concave or flat areas that meet along one edge of the boring.

Phrixichnus was previously recorded in the Pleistocene of Rhodes (Bromley and Asgaard, 1993a). It was also recognized on the Middle Miocene rocky-shores of Catalonia, Spain (Domènech et al., 2001).

***Oichnus* Bromley, 1981**

***Oichnus* isp.** (Plate 3, Figs. F, G, H)

Description: Smooth, vertical, circular to subcircular holes with axes oriented perpendicular to host substrates (gastropods and bivalves), completely or incompletely penetrative. Diameters range from 1.5–2 mm.

Remarks: *Oichnus* are generally interpreted as praedichnia of the gastropod families Naticidae and Muricidae respectively (Bromley, 1981; Pickerill and Donovan, 1998).

***Trypanites* Mägdefrau, 1932**

cf. *Trypanites* isp. (Plate 3, Fig. D)

Description: *Trypanites* is a simple, vertically to obliquely oriented boring that can curve slightly and have rounded terminations. The boring extends into the substrate from the walls of *G. torpedo* and is 1 mm diameter and up to 7 mm in length.

Remarks: *Trypanites* is generally considered to have been produced by sipunculid worms (Bromley, 1992; Pemberton et al., 1980). This trace occurs in marine carbonate deposits, including firm- to hardgrounds, pebbles and skeletal substrates (Bromley and D'Alessandro, 1987).

6. Discussion

The Miocene succession in the northeastern Aegean region indicates the first Mediterranean marine transgression affecting these areas. Frequent marine flooding occurred during the Late Miocene in the northeastern Aegean region. This period in the study area is represented by brackish to fresh water carbonates with some marine sandstone-siltstone interbeds of the Alcitepe Formation, with endolithic bivalves, endolithic clionid sponges, echiroids, polychaetes, annelids and ostracods.

Nineteen ichnotaxa producing bioerosional structures are identified in this environment: *G. torpedo*, *G. lapidicus*, *Gastrochaenolites* isp., *Phrixichnus* isp., *E.goniodes*, *E.geometrica*, *E.laquea*, *E.ovula*, *E.solaris*, *Entobia* isp., *Maeandropolydora* isp., *M.sulcans*, *M.decipiens*, *Caulostrepis taeniola*, *Caulostrepis* isp., cf. *Circolites* isp., cf. *Trypanites* isp., cf. *Conchotrema* isp., cf. *Oichnus* isp..

The foremost trace fossil in the studied area is *Gastrochaenolites*, which is a typical hardground colonizer. Two ichnospecies of *Gastrochaenolites* have been identified on the basis of the shape of the distal part of the boring. *G. lapidicus*, which displays a rounded base, and *G. torpedo*, which is larger with a sharper base. Some of the borings contain the trace-producing organism in living position, and in some the proximal part of *Gastrochaenolites* is truncated by submarine erosion. *Phrixichnus* has also been observed for the first time. *Phrixichnus* is a clavate boring resembling *Gastrochaenolites*.

Entobia is the second most dominant ichnogenus in the studied material. Five ichnospecies are identified in the study area. It is frequently observed together with *Gastrochaenolites*. The inferred polychaete annelid borings, sipunculid worms, mollusc drill holes, and duraphagous scars are also present with *Gastrochaenolites* and *Entobia*.

The rocky-shore in the study area is covered with limestone boulders, which correspond to the bioeroded surfaces of wave-cut platforms. These result from cliff retreat, which is on relatively large sub-horizontal rocky surfaces. As a stratigraphic point of view, the above mentioned bioeroded rockground surfaces reflect hard substrate marine flooding surfaces (transgressive surfaces). The hardground surface shows well-developed macro-bioerosional structures. According to previous workers (Santos et al., 2010) the hardground marine flooding surfaces (transgressive surfaces) are characterized by the presence of *Gastrochaenolites* borings, especially vertical *G. torpedo*.

Therefore, the rocky-shores at Bozcaada Island, including bioerosional structures with an abundance of the borings *Gastrochaenolites*, especially *G. torpedo*, can be interpreted as horizontal wave-cut platforms with low or no sedimentation. Moreover, the occurrences of this ichnogenus (*Gastrochaenolites*) characterize shallow water environments (Bromley, 1992; Bromley and Asgaard,

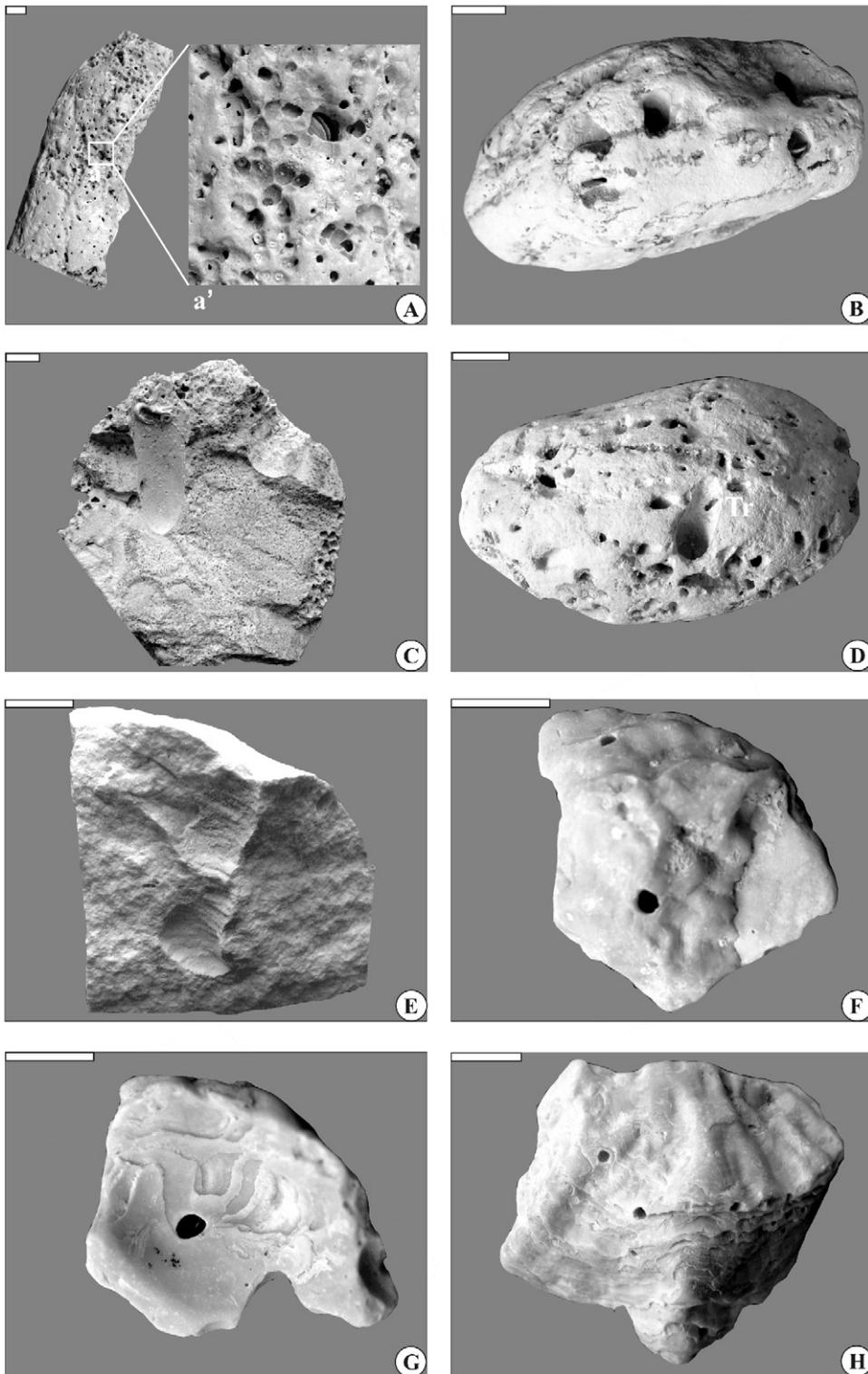


Plate 3. Images of inferred bivalve borings and drill holes-duraphagous scars from the Alçıtepe limestone boulders and shells (All scales: 1 cm). **A.** *Gastrochaenolites* isp. **B.** *Gastrochaenolites lapidicus*. **C.** *Gastrochaenolites torpedo*. **D.** *Gastrochaenolites torpedo* with *Trypanites* isp. **E.** *Phrixichnus* isp. **F.** *Oichnus* isp. **G.** *Oichnus* isp. **H.** *Oichnus* isp.

Planche 3. Images de trous et creusements supposés de bivalves-traces de duriphages dans des blocs et coquilles calcaires de la Formation d'Alçıtepe (toutes échelles = 1 cm). **A.** *Gastrochaenolites* isp. **B.** *Gastrochaenolites lapidicus*. **C.** *Gastrochaenolites torpedo*. **D.** *Gastrochaenolites torpedo* with *Trypanites* isp. **E.** *Phrixichnus* isp. **F.** *Oichnus* isp. **G.** *Oichnus* isp. **H.** *Oichnus* isp.

1993a, b), according to Bromley (1992), indicating only a few metres of water depth. Water depth is restricted to 1–2 m in the Neogene of the Mediterranean areas affected by *G. torpedo* (Kleemann, 1973). However, unpublished work has recently revealed that *Gastrochaenolites* may have occurred in several tens of meters water depth in the past. G.R. Bromley and U. Asgaard (personel communication) have observed *Gastrochaenolites* in ancient Greek marble statues from a Roman shipwreck at an approximate depth of 45 m. Also Danian *Gastrochaenolites* in Denmark seems to have inhabited deeper-water (Kleemann, 1973).

On the other hand, all endolithic organisms, especially *Entobia* borings (endolithic sponges), require low rates of sedimentation. In this context, the studied ichnoassemblage suggests a rocky-shore with a low rate of sedimentation. In addition, low sedimentation or non-deposition are revealed by bioeroded hardgrounds (Cachão et al., 2008). The bioeroded surfaces, recognized in the field on the wave-cut platform, being hardground transgressive marine flooding surfaces, are characterized by *Gastrochaenolites* borings. The assemblages found on wave-cut platforms are composites resulting from the overprinting of successive ichnocenoses that could replace each other during a transgression. As the transgression progressed and the substrate was exposed to deeper conditions, an initial bivalve-dominated community was progressively replaced by a sponge-dominated community.

7. Conclusions

1. The study of the Late Miocene rocky-shore in the Habbeli Bay area (western part of Bozcaada Island) in the Alcitepe Formation revealed an assemblage consisting of the following 19 ichnotaxa: *Oichnus* isp., cf. *Conchotrema*, *Circolites* isp., *Entobia* isp., *E. goniodes*, *E. geometrica*, *Entobia laquea*, *Entobia ovula*, *Entobia* cf. *solaris*, *Gastrochaenolites* isp., *G. torpedo*, *G. lapidicus*, *Phrixichnus* isp., *Maeandropolydora* isp., *Maeandropolydora sulcans*, *Maeandropolydora decipiens*, *Caulostrepis* isp., *Caulostrepis teniola*, and cf. *Trypanites*.

2. The ichnoassemblage associated with the limestone boulders collected from the rocky-shore is dominated by the trace of endolithic bivalves, endolithic sponges, echinoids, polychaetes, and annelids. These are marine organisms, which belong to a marine hard substrate community showing the existence of an ancient wave-cut platform.

3. *Gastrochaenolites* and *Entobia* are dominant borings in the studied locality.

4. Although bivalve borings of the ichnogenus *Gastrochaenolites* and the sponge boring *Entobia* are dominant in this region, in the shallower part, *Gastrochaenolites* is more dominant (Bromley and Asgaard, 1993a; Martinell and Domènech, 1995). As the transgression progressed and substrate was exposed to deeper conditions, an initial bivalve-dominated community was progressively replaced by a sponge-dominated community. During this period of time, bioerosion combined with physical erosion lowered the substrate, thus destroying a great part and truncating the *Gastrochaenolites* and *Entobia*.

5. *Phrixichnus* has been for the first time reported in Turkey.

6. The bioerosional assemblage found in boulders consists of small *Gastrochaenolites* and *Entobia*. They indicate that endolithic communities on unstable substrates were affected by physical factors.

7. The borings can be classified as a *Gastrochaenolites-Entobia* assemblage, which is typical of Neogene rocky-shores and belongs to the *Entobia* ichnofacies. This characterizes littoral rockground environments indicating wave-cut platforms and marine flooding surfaces (transgressive surfaces) in the stratigraphical record.

Acknowledgements

This work was supported by the Research Project 2009-08-16-01-2 of the Natural History Museum General Directorate of Mineral Research and Exploration Ankara, Turkey. The author is grateful to Prof. Dr. Richard Bromley (Copenhagen University, Denmark) for critical remarks and improvement of the manuscript, and to Prof. Dr. Ana Santos (Universidad de Huelva, Spain) who made valuable suggestions. Also acknowledged is Dr. Ceren Küçükuysal, Ayşe Demirci, and Tuncay Temiz for their technical support during the preparation of this paper, and Prof. M.C. Goncuoglu for linguistic improvements.

Finally, I wish to thank Dr. Michel Laurin (Editor) and reviewers for the constructive discussions.

References

- Brett, C.E., 1988. Paleoecology and evolution of marine hard substrate communities: an overview. *Palaios* 3, 374–378.
- Bromley, R.G., 1981. Concept in ichnotaxonomy illustrated by small round holes in shells. *Acta Geologica Hispanica* 16, 55–64.
- Bromley, R.G., 1992. Bioerosion: eating rocks for fun and profit. In: Maples, C.G. and West, R.R. (Eds.). *Trace Fossils. Short Course in Paleontology* 5, pp. 121–129.
- Bromley, R.G., 1994. The palaeoecology of bioerosion. In: Donovan, S.K. (Ed.), *The palaeobiology of trace fossils*. John Wiley and Sons, Chichester, pp. 134–154.
- Bromley, R.G., D'Alessandro, A., 1983. Bioerosion in the Pleistocene of Southern Italy: ichnogenes *Caulostrepis* and *Maeandropolydora*. *Rivista Italiana di Paleontologia e Stratigrafia* 89, 283–309.
- Bromley, R.G., D'Alessandro, A., 1984. The ichnogenes *Entobia* from the Miocene, Pliocene and Pleistocene of southern Italy. *Rivista Italiana di Paleontologia e Stratigrafia* 90, 227–296.
- Bromley, R.G., D'Alessandro, A., 1987. Bioerosion of the Pli - Pleistocene transgression of southern Italy. *Rivista Italiana di Paleontologia e Stratigrafia* 93, 379–422.
- Bromley, R.G., D'Alessandro, A., 1989. Ichological study of shallow marine endolithic sponges from the Italian coast. *Rivista Italiana di Paleontologia e Stratigrafia* 95, 279–314.
- Bromley, R.G., Asgaard, U., 1993a. Endolithic community replacement on a Pliocene rocky-coast. *Ichnos* 2, 93–116.
- Bromley, R.G., Asgaard, U., 1993b. Two bioerosion ichnofacies produced by early and late burial associated with sea level change. *Geol. Rundsch.* 82, 276–280.
- Bronn, H.G., 1837. *Lethaea geognostica*. 2. Das Kreide und Molassen - Gebirge pp. 545–1350.
- Cachão, M., Silva, C., Santos, A., Domènech, R., Martinell, J., Mayoral, E., 2008. The bioeroded megasurface of Oura (Alvare, South Portugal): implications for the Neogene stratigraphy and tectonic evolution of Southwest Iberia. *Facies* 55, 213–225.
- Clarke, J.M., 1908. The beginnings of dependent life. *Bulletin of New York State Museum* 121, 146–196.
- Çağatay, N., Görür, N., Flecker, R., Sakıncı, M., Tunçoğlu, C., Ellam, R., Krijgsman, W., Vincent, S., Dikbaş, A., 2006. Paratethyan - Mediterranean connectivity in the Sea of Marmara region (NW Turkey) during the Messinian. *Sedim. Geol.* 188–189, 171–187.

- Dermitzakis, M.D., Papanikolaou, D.J., 1981. Paleogeography and geodynamics of the Aegean region during the Neogene. *Annales Geologiques des Pays Helleniques (hors serie)* 4, 245–289.
- Domènech, R., de Gibert, J.M., Martinell, J., 2001. Ichnological features of a marine trasgression: middle miocene rocky-shores of Tarragona, Spain. *Geobios* 34, 99–107.
- Druit, C.E., 1961. Report on the petroleum prospect of Thrace, Turkey: Turkish Gulf Oil Co. (Unpublished).
- Ekdale, A.A., Bromley, R.G., Pemberton, S.G., 1984. Ichnology. The use of trace fossils in sedimentology and stratigraphy. Society of Economic Paleontologists and Mineralogists, Tulsa, short course no. 15, 317 p.
- Fisher, R., 1990a. Significado paleoecológico y geológico de perforaciones fósiles de bivalvos. *Revista de la Sociedad Mexicana de Paleontología* 3, 79–95.
- Fisher, R., 1990b. Paläoökologische Bedeutung fossiler Muschelbohrungen. *Arbeitskreis Paläontologie Hannover* 18, 1–28.
- Ghibaud, G., Grandesso, P., Massari, F., Uchman, A., 1996. Use of trace fossils in delineating sequence stratigraphic surfaces (Tertiary Venetian Basin, northeastern Italy). *Palaeogeogr. Palaeoclimat. Palaeoecol.* 120, 261–279.
- Gibert, J.M., Martinell, J., Domènech, R., 1998. *Entobia* ichnofacies in fossil rocky-shores. Lower Pliocene, northwestern Mediterranean. *Palaios* 13, 476–487.
- Göncüoğlu, M.C., 2010. Introduction to the Geology of Turkey: Geodynamic evolution of the pre-Alpine and Alpine terranes. General Directorate of Mineral Research and Exploration Monography Series, 5, 1–66. (ISBN 978-605-4075-74).
- Hohman, J.I., 1993. 6th Keck Symposium, Whitman College, pp. 236–239.
- Jia-Yu, Rong, Johnson, M.E., 1996. A stepped karst unconformity as an early Silurian rocky-shoreline in Guizhou Province (South China). *Palaeogeogr. Palaeoclimat. Palaeoecol.* 121, 115–129.
- Johnson, M.E., 1992. Ancient rocky-shores: a brief history and annotated bibliography. *Journal of Coastal Research* 8, 797–812.
- Jones, B., Pemberton, S.G., 1988. *Lithophaga* borings and their influence on the diagenesis of coral in the Pleistocene Ironshore Formation of Grand Cayman Island, British West Indies. *Palaios* 3, 3–21.
- Kelly, S.R.A., Bromley, R.G., 1984. Ichnological nomenclature of clavate borings. *Paleontology* 27, 793–807.
- Kesgin, Y., Varol, B., 2003. Gökçeada ve Bozcaada'nın Tersiyer Jeolojisi (Çanakale). *Maden Tetkik ve Arama Dergisi* 126, 49–68 (in Turkish).
- Kleemann, K.H., 1973. Der Gesteinsabbau durch Ätzmuscheln an Kalkküsten. *Oecologia* 13, 377–395.
- Kobluk, D.R., 1981a. The record of early cavity-dwelling (coelobiontic) organism in the Paleozoic. *Canadian J. Earth Sci.* 18, 181–190.
- Kobluk, D.R., 1981b. Earliest cavity-dwelling organism (coelobiontic), Lower Cambrian Poleta Formation, Nevada. *Canadian J. Earth Sci.* 18, 669–679.
- Kobluk, D.R., James, N.P., 1979. Cavity-dwelling organism in Lower Cambrian patch reefs from southern Labrador. *Lethaia* 12, 193–218.
- Kobluk, D.R., James, N.P., Pemberton, S.G., 1978. Initial diversification of macroboring ichnofossils and exploitation of macroboring niche in the Lower Paleozoic. *Paleobiology* 4, 163–170.
- Leymerie, M.A., 1842. Suite des mémoires sur le terrain Crétacé du département de l'Aube. *Mem. Soc. Geol. France* 5, 1–34.
- Mägdefrau, K., 1932. Über einige Bohrgänge aus dem Unteren Muschelkalk von Jena. *Paläontologische Zeitschrift* 14, 150–160.
- Martinell, J., 1989. Interaccion organismos/sustrato duro: la bioerosion y sus aplicaciones. In: Aguirre, E. (Ed.), *Paleontologia. Colección "Nuevas Tendencias"* 10, pp.205–222.
- Martinell, J., Domènech, R., 1995. Bioerosive structures on the Pliocene rocky-shores of Catalonia (Spain). *Revista Española de Paleontología* 10, 37–44.
- Mikuláš, R., 1992. Early Cretaceous borings from Štramberk (Czechoslovakia). *Časopispro Mineralogii a Geologii* 37, 297–312.
- Neumann, A.C., 1966. Observations on coastal erosion in Bermuda and measurements of the boring rate of the sponge *Cliona lampa*. *Lymnology and Oceanography* 11, 92–108.
- Önem, Y., 1974. Gelibolu ve Çanakale dolaylarının jeolojisi. TPAO Raporu: 877 (Unpublished) (in Turkish).
- Pemberton, S.G., Kobluk, D.R., Yeo, R.K., Risk, M.J., 1980. The boring *Trypanites* at the Silurian-Devonian disconformity in southern Ontario. *J. Paleontol.* 54, 1258–1266.
- Pickerill, R.K., Donovan, S.K., 1998. Ichnology of the Pliocene Bowden shell bed, southeast Jamaica. *Contr. Tert. Quat. Geol.* 35, 161–175.
- Radwański, A., 1969. Lower Tortonian transgression onto the southern slopes of the Holy Cross Mts. *Acta. Geologica Polonica* 19, 164–177.
- Radwański, A., 1970. Dependence of rock borers and burrowers on the environmental conditions within the Tortonian littoral zone of Southern Poland. In: Crimes, T.P., Harper, J.C. (Eds.), *Trace Fossils 2: Geol. J. Special Issues* 9, pp.227–264.
- Sakıncı, M., Yaltrak, C., 1997. Marine Pleistocene deposits and palaeogeography along the Southern Thrace coastal area. *Maden Tetkik Arama Dergisi* 119, 43–62 (in Turkish).
- Sakıncı, M., Yaltrak, C., 2005. Messinian crisis: what happened around the northeastern Aegean? *Marine Geol.* 221, 423–436.
- Sakıncı, M., Yaltrak, C., Oktay, F.Y., 1999. Palaeogeographical evolution of the Thrace Neogene Basin and the Tethys-Paratethys relations at northwestern Turkey (Thrace). *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 153, 17–40.
- Santos, A., Mayoral, E., Silva, C.M., Cachão, M., Domènech, R., Martinell, J., 2008. Trace fossil assemblages on Miocene rocky-shores of southern Iberia. In: Wishak, M., Tapanila, L. (Eds.), *Current developments in bioerosion*. Springer, Berlin, Heidelberg, New York, pp. 431–450.
- Santos, A., Mayoral, E., Silva, C.M., Kullberg, C.J., 2010. Trypanites ichnofacies: palaeoenvironmental and tectonic implications. A case study from the Miocene disconformity at Foz da Fonte (Lower Tagus Basin, Portugal). *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 292, 35–43.
- Siyak, M., Bürkan, K.A., Okay, A.I., 1989. Biga ve Gelibolu yarımadalarının Tersiyer jeolojisi ve hidrokarbon olanakları. *Türkiye Petrol Jeologları Derneği Bülteni* 3, 183–199 (in Turkish).
- Taylor, P.D., Wilson, M.A., 2002. A new terminology for marine organisms inhabiting hard substrates. *Palaios* 17, 522–525.
- Taylor, P.D., Wilson, M.A., 2003. Palaeoecology and evolution of marine hard substrate communities. *Earth Sci. Rev.* 62, 1–103.
- Teichert, C., 1945. Parasitic worms in Permian brachiopod and pelecypod shells in Western Australia. *Am. J. Sci.* 243, 197–209.
- Temel, R.Ö., Çiftçi, N.B., 2002. Gelibolu Yarımadası, Gökçeada ve Bozcaada Tersiyer çökellerinin stratigrafisi ve ortamsal özellikleri. *Türkiye Petrol Jeologları Derneği Bülteni* 14, 17–40 (in Turkish).
- Uchman, A., Demircan, H., Tokar, V., Derman, S., Sevim, S., Szulc, J., 2002. Relative sea level changes recorded in borings from a Miocene rocky-shore of the Mut Basin, southern Turkey. *Annales Societatis Geologorum Poloniae* 72, 263–270.
- Voigt, E., 1965. Über parasitische Polychaeten in Kreide-Austern sowie einige andere in Muschelschalen bohrende Würmer. *Paläontologische Zeitschrift* 39, 193–211.
- Voigt, E., 1971. Fremdkulturen an Steinkernen von Polychaeten-Bohrungen aus der Maastrichter Tuffkreide. *Paläontologische Zeitschrift* 45, 144–153.
- Voigt, E., 1975. Tunnelbaue rezenter und fossiler Phoronidea. *Paläontologische Zeitschrift* 49, 135–167.
- Warme, J.E., 1975. Borings as trace fossils and the processes of marine bioerosion. In: Frey, R.W. (Ed.), *The Study of Trace Fossils*. Springer-Verlag, New York, pp. 181–227.
- Watkins, R., 1990. Palaeoecology of a Pliocene rocky-shoreline, Salton Trough Region, California. *Palaios* 5, 167–175.
- Wisshak, M., Gektidis, M., Freiwald, A., Lundäl, T., 2005. Bioerosion along a bathymetric gradient in a cold temperate setting (Kosterfjord, SW Sweden): an experimental study. *Facies* 51, 93–117.
- Yaltrak, C., 1996. Stratigraphical and sedimentological properties of the Pleistocene marine depocentres of the southern Thrace coast. Msc. Thesis. Institute of Marine Sciences and Management, Istanbul University 48 p. (In Turkish).
- Yaltrak, C., Alpar, B., 2002. Kinematics and evolution of the northern branch of the North Anatolian fault (Ganos Fault) between the Sea of Marmara and the Gulf of Saros. *Marine Geol.* 190, 351–366.
- Yaltrak, C., Alpar, B., Yüce, H., 1998. Tectonic elements controlling the evolution of the Gulf of Saros (Northeastern Aegean Sea). *Tectonophysics* 300, 227–248.
- Yaltrak, C., Sakıncı, C., Oktay, F.Y., 2000. Kuzey Anadolu Fayı'nın Kuzey Kolunun (Ganos Fayı) Marmara ve Saroz Körfezi Arasındaki Kinematığı ve Evrimi. In: Tatar, O., Kavak, K.Ş., Özden, S. (Eds.), *ATAG-III Workshop Kitabı*, pp. 77–89 (In Turkish).
- Yaltrak, C., Sakıncı, M., Aksu, A.E., Hiscott, R.N., Galle, B., Ulgen, U.B., 2002. Late Pleistocene uplift history along the southwestern Marmara Sea determined from raised coastal deposits and global sea level variations. *Marine Geol., Special Issue on the Sea of Marmara* 190 (1/2), 283–306.