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MAASTRICHTIAN - SELANDIAN PLANKTONIC FORAMINIFERA BIOSTRATIGRAPHY AND PALAEOECOLOGICAL INTERPRETATION OF AKVEREN FORMATION IN BARTIN AREA (WESTERN BLACK SEA, TURKEY)

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Research Article

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ABSTRACT

The changes in planktonic foraminifera assemblages were studied in two sections composed of clayey limestone, limestone, claystone and marl in the Akveren Formation in the Bartın Province. *Pseudoguembelina palpebra*, *Racemiguembelina fructifera*, *Abathomphalus mayaroensis* and *Pseudoguembelina hariaensis* biozones in the Maastrichtian and *Parvularugoglobigerina eugubina*, *Glomobanomalina compressa*/*Praemurica uncinata*, *Praemurica uncinata* and *Morozovella angulata* biozones in the Paleocene were determined using planktonic genus and species identified in these sections. Paleocological interpretations in this study were completed by using the relative abundances of paleoecological species identified in these biozones. The abundance of *Rugoglobigerina* spp., *Heterohelix globulosa*, *Pseudoguembelina* spp., species that tolerated changes in sea water temperature and nutrition, increased in the Maastrichtian. In the uppermost Maastrichtian, high abundance of *Racemiguembelina fructifera*, *Heterohelix globulosa* and *Rugoglobigerina* spp. show oligotrophic and warmer environmental conditions in this study. The abundance of globotruncanids are unstable due to environmental changes. The new species have smaller sizes and globular chambers emerged in the Paleocene. High abundance of *Subbotina triloculinoides* and *Parasubbotina pseudobulloides* show mesotrophic and cooler environmental conditions in the Danian. Morozovellid taxa started to dominate in the Selandian and environmental conditions changed from mesotrophic to oligotrophic.

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1. Introduction

The temperature in the Late Cretaceous from Cenomanian to Maastrichtian was higher than today as determined by stable isotope studies and global climate models (Huber et al., 1995, 2002; Clarke and Jenkyns, 1999; DeConto et al., 1999; Wilson and Norris, 2001; Norris et al., 2002; Wilson et al., 2002; Bice et al., 2006; Forster et al., 2007; Bornemann et al., 2008; Friedrich et al., 2008; Shirazi et al., 2013; Kaya-Özer, 2014).

The Cretaceous climate started to cool in the Campanian and Maastrichtian (Huber et al., 1995, 2002). While the decrease reached a maximum level in the Maastrichtian (Clarke and Jenkyns, 1999; Huber et al., 2002; Miller et al., 2005), a sudden increase in temperature occurred at the end of this period (Li and Keller, 1998a,b) and surface water temperatures increased by 2-3 °C between ~65.45 and 65.1 Ma before the Cretaceous–Paleogene boundary (K–Pg) (Li and Keller 1998a,b; Barrera and Savin,

1999). Global cooling continued in the last 100 kyr of the Maastrichtian and sea surface water temperatures decreased (Thibault and Gardin, 2007).

Climatic changes caused variations in the biota of planktonic foraminifera together with most fauna (Abramovich and Keller, 2003). An important biotic event that occurred in this period is a decrease in planktonic foraminifera species richness at about 66 Ma in the South Atlantic and Tethys basins, coincident with the end-Maastrichtian global warming (Li and Keller, 1998a, b). This was also proven by oxygen isotope studies carried out on deep-sea drillings in the Atlantic, Pacific and Indian Oceans (Zachos et al., 1985, 1989; D'Hondt and Lindinger, 1994; Corfield and Norris, 1996; Barrera et al., 1997; Li and Keller, 1998a,b; Abramovich and Keller, 2003; Friedrich et al., 2012) and in the Boreal Realm (Friedrich et al., 2008).

Keller et al. (2002) noticed the faunal decline crested at the K/Pg boundary with the mass extinction

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of tropical-subtropical planktonic foraminiferal species whereas the cosmopolitan and ecologically generalist species survived into the Danian. At first, tropical species with complex, large and ornamental shells (*globotruncanids*, *racemiguembelinids*, *planoglobulinids*) and then smaller subtropical species (*pseudotextularids*, *rugoglobigerinids*) were affected during the extinction at the K/Pg boundary (Keller, 1996)

There are lots of paleoecological studies based on stable oxygen and carbon isotopes and changes in planktonic foraminifera assemblages in Maastrichtian and Paleocene sediments which were collected from deep sea drillings and several locations worldwide (Huber et al., 1995; Li and Keller, 1998a). The aim of this study is to identify planktonic foraminifera biostratigraphy in the Akveren Formation and determine the paleoecological conditions based on changes in planktonic foraminifera assemblages between Maastrichtian and lower Paleocene period using samples from Bartın Province (Western Black Sea) in Turkey. The biostratigraphic and paleoenvironmental interpretations of the composition of the planktonic foraminifers were also discussed.

2. Geological Setting and Previous Studies

The study area is in the Western Black Sea Basin in the Pontides, one of the tectonic plates in Turkey, which is a part of the Alpine Belt. There are several geological studies about Bartın province and its surrounding. The first studies in the region were about its petroleum potential (Badgley, 1959; Saner, 1981; Gedik and Korkmaz, 1984; Robinson et al., 1996; Görür and Tüysüz, 1997). There are also several studies investigating its geological and tectonic characteristics. The geological map of Cide-Kurucaşile area was drawn by Akyol et al. (1974). Geological structure, tectonic, petrographical and petrological characteristics of the region from Jurassic to Quaternary were investigated by Saner (1980), Gedik et al. (1983), Barka et al. (1983), Aydın et al. (1986), Derman et al. (1995), Derman (2002) and Tüysüz et al. (2012).

Şengör (1982) suggested that the Western Black Sea basin was opened as a back arc basin in relation to closure of the Tethys Ocean. Görür (1988) mentioned the Western Black Sea Basin as related to post rift thermal subsidence (Görür, 1988). Özçelik and Çaptuğ (1990) examined the tectonic characteristics

of the region and emphasized that sedimentation was continuous in upper Cretaceous/Tertiary units. In addition, they pointed out that the Akveren Formation is of Maastrichtian-early Paleocene age, comprises marls, calciturbiditic limestones, micrites and shale.

Akman (1992) studied the stratigraphic units between Lutetian and Permian, and named the various sediments, volcano-sedimentary, and volcanic rocks that outcrop in the region. Tüysüz (1993) investigated the Pontides in detail, classified its tectonic sections and suggested a geological evolution model. Sunal and Tüysüz (2001) studied the stratigraphy and tectonics of the region. Tüysüz (2002) stated that deposition was accompanied by strong volcanism in the Cenomanian-Maastrichtian period and this volcanism that was common on the Black Sea coast began at the end of the Campanian and ended in the Maastrichtian.

Paleontological studies completed in the study area and its surrounding are important. Dizer and Meriç (1981) defined the planktonic foraminifera zones from the upper Cretaceous to Paleocene in Gebze, Akçakoca, Devrek and Bartın areas in Northwest Anatolia. They stated that the sea had a deep sea characteristic in Bartın in the upper Cretaceous and Paleocene. Varol (1983) discussed the late Cretaceous-Paleocene calcareous nanofossils from the Kokaksu Section. Sirel (1991) defined a new species (*Cideina* n.gen.) in his study in the Cide region. Özkan-Altınar and Özcan (1999) examined the benthic and planktonic foraminifera in the units from late Cretaceous to early Tertiary in the forearc basins located in north, northwest and central Anatolia.

Şener (2007) identified *Neotrocholinitid* and *Orbitolinid* species that were benthic foraminifera in the İnaltı Formation outcropping in Amasra. Three new *Neotrocholinita* species (*Neotrocholinita amasraensis*, *N. sireli*, *N. bartinensis*) were identified, the age of the formation in the study area was defined as Cimmerian-Aptian based on these identified species.

3. Stratigraphy

The oldest unit in the area consists of Amasra Group rocks, belonging to the Yemişliçay Upper Group (Tüysüz et al., 2004). The Amasra Group, deposited in the late Santonian and Campanian, begins with a marine succession that overlaps the older units (Tüysüz et al., 2004). At the beginning of this period, pelagic limestones were deposited and then volcanics

and pyroclastics (Kazpınar Formation) developed (Tüysüz et al., 2004).

3.1. Kazpınar Formation

The name of the formation was first used by Tokay (1954/1955). This unit was studied as the Yemişliçay formation (Ketin and Gümüş, 1963), Kuruçayı formation (Akyol et al., 1974), Lüneran formation (Kaya et al., 1982/1983), Cambu formation (Şahintürk and Özçelik, 1983; Tüysüz et al., 1997) and Dinlence formation (Akman, 1992) in previous studies. Kazpınar formation widely consists of andesitic lavas, pyroclastics, volcanic sandstones, siltstones, marls and lava blocks (Tüysüz et al., 2004).

Kazpınar formation indicates volcanic activity in the deep sea environment (Akman, 1992). Turbiditic units were deposited when the volcanism lost its effect (Akman, 1992). The age of the formation was defined as Campanian according to its stratigraphic position (Tüysüz et al., 2004). In another study, it was dated as early Campanian due to genus and species of planktonic foraminifera and nanofossils that were identified in carbonate levels in this formation (Kaya-Özer, 2009).

3.2. Akveren Formation

Akveren Formation overlaps the Yemişliçay group rocks consisting of volcanic and volcanoclastic rocks (Gedik et al., 2005). The Akveren Formation was first described by Gayle (1959) as layers of clayey limestone then formally defined by Ketin and Gümüş (1963) as widespread along the western Black Sea coast.

Akveren Formation was the subject of several studies. Dobruçalı (1985) and Sarıca (1993) examined planktonic foraminifera fauna and stratigraphic position of the Akveren Formation. Kırıcı (1998) stated that the Akveren Formation includes a rich foraminifera assemblage that consists of planktonic foraminifera, small benthic foraminifera and large benthic foraminifera in Maastrichtian and dated it as late Cretaceous-Paleocene. In addition, she pointed out that it was dominated by deepening intensity flows and represented a calciturbiditic succession that reflected deposition in the open shelf environment.

Aydın (2005) studied nanoplankton biostratigraphy in detail in the units of Cretaceous-Paleogene age including the Akveren Formation northwest of İzmit. In this study, the Cretaceous-Tertiary

boundary was determined to be continuous. Güray (2006) determined the Campanian-Maastrichtian boundary by using planktonic foraminifera. While the Campanian-Maastrichtian boundary was determined as the boundary of zones of *Pseudotextularia elegans* and *Planoglobulina acervuloides* in her study, the Cretaceous/Tertiary boundary was determined by the extinction of upper Cretaceous fossils.

The Akveren Formation is initially characterized at the base by white-gray, medium to thickly bedded sandstones and greenish-gray marls that change to white-cream, thinly bedded micritic limestones and clayey limestones in the upper levels. The formation ends with white-beige, locally siliceous, thin to medium bedded limestones, claystones, and marl alternations and gradually passes into the Atbaşı Formation consisting of calcareous mudstones (Akman, 1992, Figure 1).

Thickness of the Akveren Formation was indicated as 390 m by Akyol et al. (1974) and 590 m by Akman (1992). The age of the formation was indicated as Campanian-Maastrichtian by Akman (1992), Maastrichtian by Ketin and Gümüş (1963) and Maastrichtian-Paleocene by Gedik and Korkmaz (1984). Kaya-Özer (2009), Kaya-Özer and Toker (2012) pointed out that the age of the Akveren Formation age is Campanian-Selandian based on planktonic foraminifera and nannofossil biostratigraphy.

3.3. Atbaşı Formation

The Atbaşı formation was first named by Ketin and Gümüş (1963) and different names have been used in some studies (Akyol et al., 1974; Akman, 1992). The unit consists of green and purple thin bedded, conchoidal fracture, fossiliferous siltstones, claystones and marls. Thin to medium bedded, green, greenish, gray sandstone layers and rarely white colored, thin bedded limestones are found in the unit. According to Gedik and Korkmaz (1984), its thickness is 537 m Akyol et al. (1974) named this unit the Akgüney Formation and indicated its thickness as 260 m. The age of the unit was determined as Paleocene-Early Eocene (Ketin and Gümüş, 1963), Paleocene (Akyol et al., 1974), early Eocene (Gedik and Korkmaz, 1984) and Paleocene (Tüysüz et al., 1997), in various studies. Kaya-Özer and Çakır (2015) pointed out the age of the formation was Selandian-Ypresian according to detailed planktonic foraminifera biostratigraphy.

4. Material and Method

This study is based on 37 samples from two sections in the Akveren Formation. The study area lies in the Bartın area on the Black Sea coast and is shown on the 1:25000 Zonguldak E28-c2 map (Figure 1).

The samples of planktonic foraminifera were disaggregated in 10% concentrated hydrogen peroxide soaked with water and washed through >63 µm sieve until clean foraminiferal residues were recovered. The washed samples were dried at room temperature. From each Cretaceous sample, about 300 planktonic foraminifera were picked from random sample splits (Abramovich and Keller, 2003). All Paleocene planktonic foraminifera species were picked from the samples, because they are rare. The planktonic foraminifera species were identified under a stereo

microscope (Leica zoom 2000). The important species were photographed with a scanning electron microscope (SEM-JEOL JSM-6490LV)) at the Turkish Petroleum Company (TPAO) and are presented in the plate.

Mesozoic taxonomic concepts for planktonic foraminifera identification were applied in this study according to Robaszynski et al. (1984), Caron (1985), Robaszynski and Caron (1995), Huber et al. (2008), Petrizzo et al. (2011). Cenozoic taxonomic concepts for planktonic foraminifera identification were used according to Toumarkine and Luterbacher (1985), Berggren and Norris (1997) and Olsson et al. (1999).

The percentages of planktonic foraminifera species were calculated for paleoecological interpretation and these percentages were counted according to the

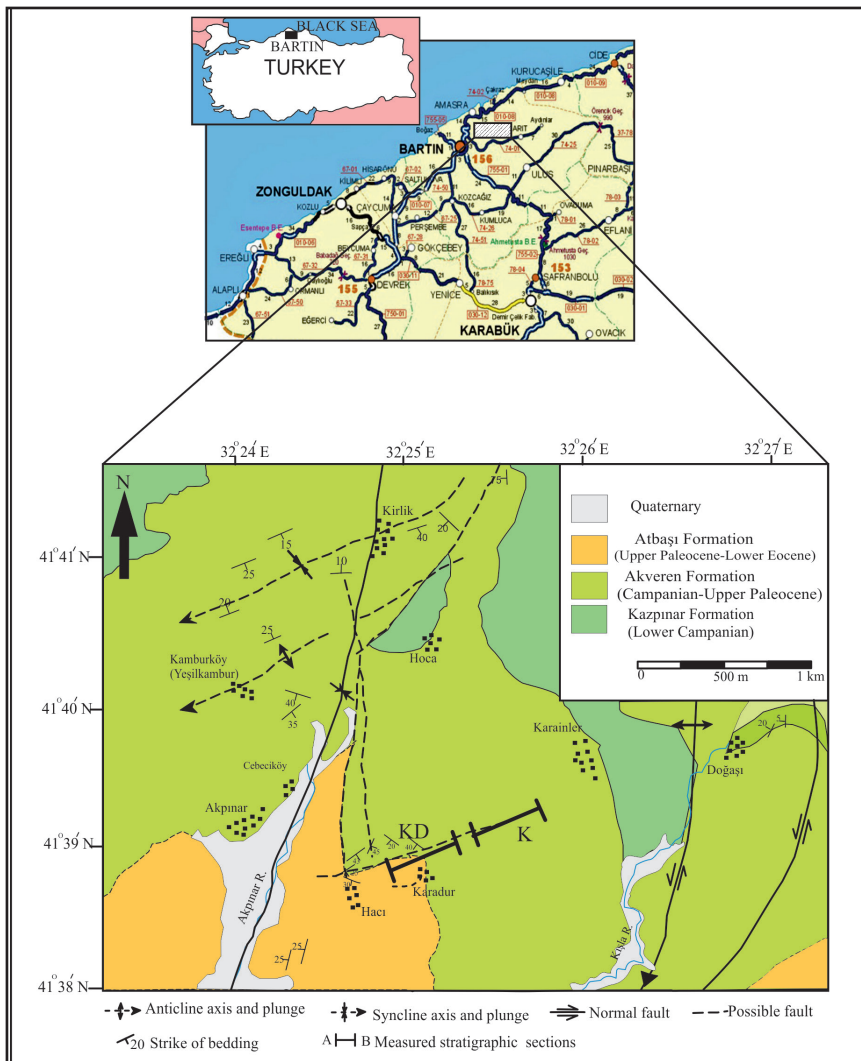


Figure 1- Location and geological map of the study area with locations of the measured sections (modified from Akman, 1992).

relative abundance of each species in the samples. The percentage abundances (%) of planktonic foraminifera are listed in table 1 and table 2.

5. Results

5.1. Stratigraphic Section

The detailed planktonic foraminiferal biostratigraphy were studied in the Karainler and Karadur areas around Bartın (Figure 1).

Karainler stratigraphic section

Measured in west of Karainler Village in the Akveren Formation, the Karainler section begins

at the coordinates 41°39'25"- 32°25'80" and ends at 41°39'15"- 32°25'50". It consists of an alternation of pink, cream-colored, thin to medium bedded clayey limestones, cream-colored, thin-bedded claystones and gray conchoidal marl at the base of the Akveren Formation (Figure 2). It passes into an alternation of gray, beige, thin to medium bedded, clayey limestones, white, thin bedded claystones and gray, green laminated marl in the upper level of the Akveren Formation. The section ends with an alternation of gray marl and thin to medium bedded limestones that include hard, partly silicified and echinoid fossils (Figure 2, 3). Total thickness of the section measured in the Akveren Formation is 140 m and the sampling intervals varied every 3-10 m in this section (Figure 3).

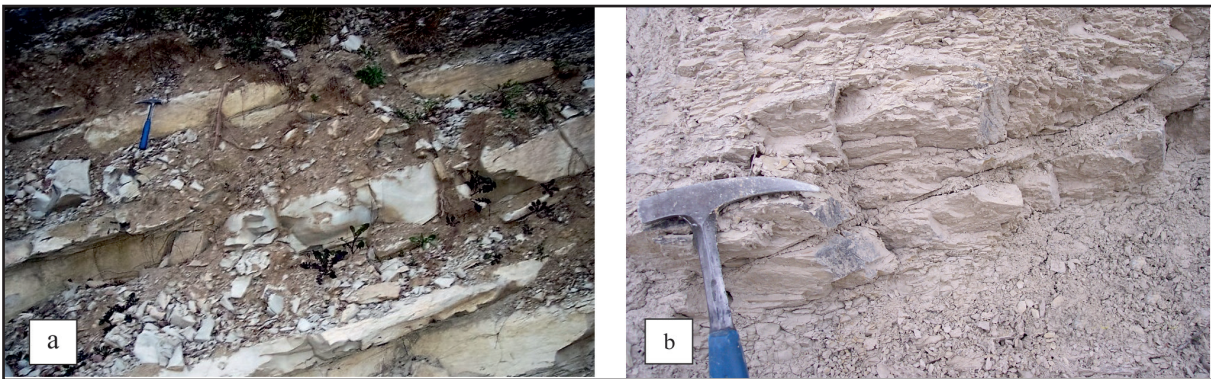


Figure 2- Field photographs of the sedimentary rocks of the Karainler stratigraphic section. (a). Alternating silicified limestones and marl in the upper Maastrichtian of the Akveren Formation, (b). interbedded gray green marls and cream clayey limestone in the lower Maastrichtian of the Akveren Formation.

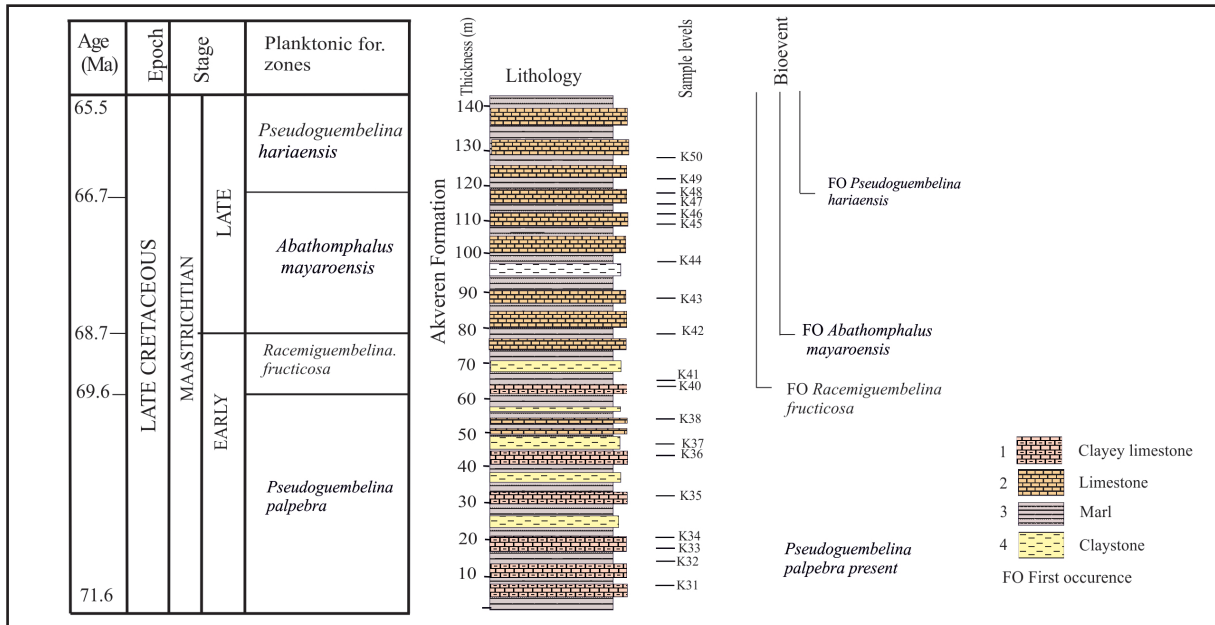


Figure 3- Karainler measured stratigraphic section showing lithostratigraphy, biostratigraphy and main planktonic foraminiferal bioevents from the Bartın area.

Karadur stratigraphic section

Measured around Karadur Village, the Karadur section begins at the coordinates 41°39'15"- 32°25'45" and ends at 41°38'80"- 32°24'95". It consists of an alternation of cream-colored, thin to medium bedded limestones, claystones and green, beige marl at the base in the Akveren Formation (Figure 4). The section continues as an alternation of partly silicified, thin to medium bedded limestones and laminated marl in the upper level of the Akveren Formation (Figure 4), it gradually passes into the Atbaşı Formation consisting of purple, green, thin to medium bedded conchoidal fracture marl in the uppermost levels (Figure 5). Total thickness of Karadur section is 23 m and the sampling intervals varied from 1-5 m in this section (Figure 5).

In the study area, the K/Pg boundary is located within the Akveren Formation. However, during the

sampling process, this boundary was missed and could not be identified.

5.2. Biostratigraphy

The standard upper Cretaceous biozonation scheme of Huber et al. (2008) and standard Paleocene biozonation scheme of Wade et al. (2011) were applied to the planktonic foraminifera data (Figure 3, 5). The important planktonic foraminifera species were shown in the plate I, II.

5.2.1. Maastrichtian Planktonic Foraminifera Zones

Pseudoguembelina palpebra Partial-range Zone: Biostratigraphic interval from the first occurrence (FO) of *Pseudoguembelina palpebra* to the FO of *Racemiguembelina fructicosa*.

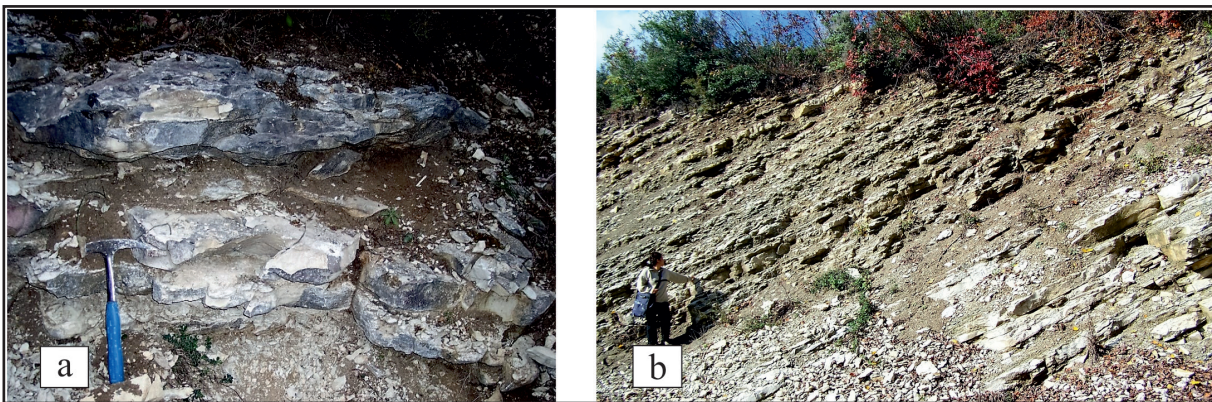


Figure 4- Field photographs of the sedimentary rocks of the Karadur stratigraphic section. (a). interbedded graygreen marls and cream silicified micritic limestones in the upper Maastrichtian of the Akveren Formation, (b). field view of Karadur stratigraphic section.

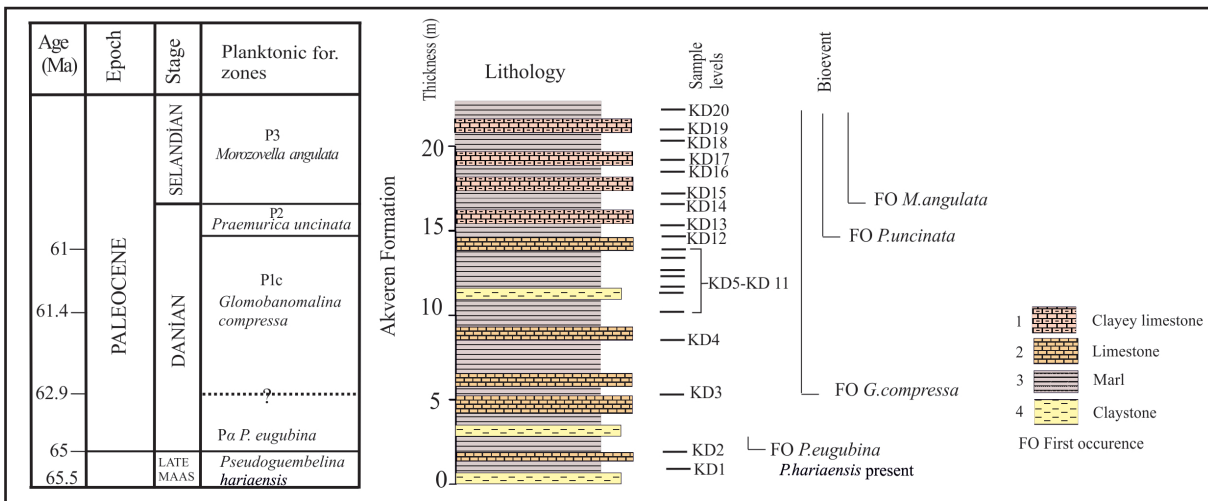


Figure 5- Karadur measured stratigraphic section showing lithostratigraphy, biostratigraphy and main planktonic foraminiferal bioevents from the Bartın area.

Author: Huber et al., 2008

Age: late Campanian-early Maastrichtian (Base 71.64 Ma, top 69.62 Ma; Huber et al. 2008).

The dominant planktonic foraminiferal species recorded in this zone include *Contusotruncana contusa*, *C. fornicata*, *C. patelliformis*, *C. plicata*, *C. plummerea*, *C. walfischensis*, *Gansserina gansseri*, *Globigerinelloides ultramicrus*, *Globotruncana arca*, *G. aegyptiaca*, *G. bulloides*, *G. falsostuarti*, *G. insignis*, *G. linneiana*, *G. mariei*, *Globotruncanella havanensis*, *G. minuta*, *G. petaloidea*, *Globotruncanita stuarti*, *G. stuartiformis*, *Heterohelix globulosa*, *Planoglobulina acervulinoides*, *P. multicamerata*, *Pseudoguembelina costulata*, *P. palpebra*, *Pseudotextularia elegans*, *Rugoglobigerina hexacamerata*, *R. macrocephala*, *R. rugosa*, *R. rotundata* (Table 1).

This zone has been identified in samples K31–K38 from the base of the Karainler stratigraphic section of the Akveren Formation (Table 1, Figure 3).

Racemiguembelina fructicosa Partial-range Zone: Biostratigraphic interval from the FO of *Racemiguembelina fructicosa* to the FO of *Abathomphalus mayaroensis*.

Author: Huber et al., 2008 (modified by Robaszynski and Caron 1995; Li and Keller 1998a, b).

Age: early Maastrichtian (Base 69.62 Ma, top 68.72 Ma; Huber et al., 2008).

The dominant planktonic foraminiferal species recorded in this zone include *Contusotruncana contusa*, *C. plicata*, *C. plummerea*, *C. walfischensis*, *Gansserina gansseri*, *Globigerinelloides ultramicrus*, *Globotruncana arca*, *G. aegyptiaca*, *G. falsostuarti*, *G. insignis*, *G. mariei*, *Globotruncanella havanensis*, *G. petaloidea*, *Globotruncanita falsocalcarata*, *G. stuarti*, *G. stuartiformis*, *Heterohelix globulosa*, *Planoglobulina acervulinoides*, *P. multicamerata*, *Pseudoguembelina costulata*, *P. palpebra*, *Pseudotextularia elegans*, *Racemiguembelina fructicosa*, *Rugoglobigerina hexacamerata*, *R. macrocephala*, *R. rugosa*, *R. Rotundata*.

This zone has been identified in samples K40 and K41 of the Karainler section (Table 1, Figure 3).

Abathomphalus mayaroensis Partial-range Zone: Biostratigraphic interval from the FO of *Abathomphalus mayaroensis* to the FO of *Pseudoguembelina hariaensis*.

Author: Huber et al., 2008, modified by Robaszynski and Caron (1995).

Age: late Maastrichtian (Base 68.72 Ma, top 66.78 Ma; Huber et al., 2008).

The dominant planktonic foraminiferal species recorded in this zone include *Abathomphalus mayaroensis*, *Contusotruncana contusa*, *C. patelliformis*, *C. plicata*, *C. plummerea*, *C. walfischensis*, *Gansserina gansseri*, *Globigerinelloides ultramicrus*, *Globotruncana arca*, *G. aegyptiaca*, *G. bulloides*, *G. falsostuarti*, *G. insignis*, *G. mariei*, *Globotruncanella havanensis*, *G. minuta*, *G. petaloidea*, *Globotruncanita falsocalcarata*, *G. stuarti*, *G. stuartiformis*, *Heterohelix globulosa*, *Planoglobulina acervulinoides*, *P. multicamerata*, *Pseudoguembelina costulata*, *P. palpebra*, *Pseudotextularia elegans*, *Racemiguembelina fructicosa*, *Rugoglobigerina hexacamerata*, *R. macrocephala*, *R. rugosa*, *R. rotundata*.

This zone has been identified in samples K42–K47 of the Karainler section of the Akveren Formation (Table 1, Figure 3).

Pseudoguembelina hariaensis Partial-range Zone: Biostratigraphic interval from the FO of the nominate species to the extinction of most Cretaceous planktonic foraminifera at the Cretaceous-Paleogene boundary.

Author: Robaszynski and Caron (1995).

Age: latest Maastrichtian (Base 66.78 Ma, top 65.50 Ma; Huber et al., 2008).

Toward the end of the Maastrichtian, globotruncanids decreased and high abundances of biserial heterohelicids, pseudoguembelinids, and rugoglobigerinids are recorded. The most characteristic elements recorded in this zone are *Abathomphalus mayaroensis*, *Contusotruncana contusa*, *C. plicata*, *C. plummerea*, *C. walfischensis*, *Gansserina gansseri*, *Globigerinelloides ultramicrus*, *Globotruncana arca*, *G. aegyptiaca*, *G. falsostuarti*, *G. mariei*, *Globotruncanella havanensis*, *G. petaloidea*,

Table 1- Percentage abundances (%) of planktonic foraminifera are represented against sample position of the Karainler stratigraphic section in the Maastrichtian.

Maastrichtian													Stage					
<i>P.palpebra</i>			<i>R.fructicosa</i>	<i>A.mayaroensis</i>					<i>Phariaensis</i>			Planktonic Foraminifera Zones						
K31	K33	K34	K35	K36	K38	K40	K41	K42	K43	K44	K45	K46	K47	K48	K49	K50	SAMPLES	
0	0	0	0	0	3.5	0	0	0.1	0	0	0	0	0.1	0	0	0	<i>Abathomphalus mayaroensis</i>	
0	0.4	0.3	0	0	0	0	0.3	1.1	0	0	0	0.2	0.2	0	0.1	0	<i>Contusotruncana contusa</i>	
17	5.3	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	<i>Contusotruncana fornicata</i>	
0.6	0	0.1	0	0	0	0	0	0.1	0	0.1	0.1	0.1	0.1	0	0	0	<i>Contusotruncana patelliformis</i>	
2.1	0.9	0.1	0	0	0	0.1	0.2	0.4	0.5	0.5	0.2	0.2	0.1	0.1	0.1	0.1	<i>Contusotruncana plicata</i>	
0	0	0.6	0	0	5.6	0.1	0.6	0.6	0	0.1	0.5	0.3	0.2	0.3	0.1	0.1	<i>Contusotruncana plummerea</i>	
0	0	0.2	0	0	14	0	0.3	0.2	0	0.3	0.3	0.2	0.2	0.1	0.1	0	<i>Contusotruncana walfischensis</i>	
0.6	1.1	0.3	0	0	2.1	0	0.2	0.3	0	0.9	0.2	0.1	0.2	0	0.1	0	<i>Gansserina gansseri</i>	
2.5	0.7	0	0	0	0	0	5	0.3	1.1	1.1	3.5	1.1	1.5	0	0	0	<i>Globigerinelloides ultramicrus</i>	
3.2	14	0.5	0.3	0.1	9.2	0.5	0	0.3	0.1	0.3	0.1	0.1	0.1	0.2	0.1	0.1	<i>Globotruncana aegyptiaca</i>	
13	8.8	1.5	1.4	1.2	7	0.4	1.7	1.5	1.2	0	1.3	1.2	1.5	0.8	0.6	0.5	<i>Globotruncana arca</i>	
0	0	0	0.1	0	0	0	0	0.1	0.1	0	0	0.1	0	0	0	0	<i>Globotruncana bulloides</i>	
0	0	0.1	0.3	0.2	1.4	0.2	0.1	0.1	0.2	0.2	0.1	0.2	0.1	0.1	0.1	0.1	<i>Globotruncana falsostuarti</i>	
0	0.4	0	0	0.1	0	0.5	0.2	0.3	0	0	0.1	0.1	0	0	0	0	<i>Globotruncana insignis</i>	
16	3.5	1.2	2.5	3.6	0	0	0	0	0	0	0	0	0	0	0	0	<i>Globotruncana linneiana</i>	
2.5	1.6	0	0.3	0.3	3.5	0.3	0	0	0	0	0	0.4	0.3	0.5	0.2	0.2	<i>Globotruncana mariei</i>	
0	5.3	0.5	8	17	0	2.9	3.1	2.2	2.7	4.9	1.9	2.2	0.2	0.1	0.2	0.1	<i>Globotruncanella havanensis</i>	
4.8	2.7	1.2	1.5	1.7	0	0	0	0	0	0	0	0.1	0.1	0	0	0	<i>Globotruncanella minuta</i>	
3	1.6	1.5	2.3	1.9	0	3.8	12	13	14	11	11	10	9	16	17	16	<i>Globotruncanella petaloidea</i>	
0	0	0	0	0	0	0.1	0.8	0	0	0.6	0.1	0.1	0.1	0.6	0.3	0.5	<i>Globotruncanita falsocalcarata</i>	
0.4	0	0.1	0	0.1	0	0.2	0.6	0.4	0.3	0.2	0.4	0.6	1.2	0.1	0.1	0.1	<i>Globotruncanita stuarti</i>	
1.3	0.5	0.4	1.1	0.2	0	1.1	0.8	0.8	0.5	0.4	0.8	0.4	0	0.1	0.2	0.1	<i>Globotruncanita stuartiformis</i>	
4.7	2.1	14	10	16	4.2	6.1	6.7	6.7	7.3	6.5	6	4.5	9.4	13	13	16	<i>Heterohelix globulosa</i>	
0.6	0.2	0.1	0	0.3	0	2.3	1.1	0.9	0	1.6	1	1.5	2.5	3.4	3.9	2.7	<i>Planoglobulina acervulinoides</i>	
0.2	0	0.1	0	0	0	0.5	0.9	0.5	0.6	0.9	0.7	0.5	0.2	1.3	0.8	1.3	<i>Planoglobulina multicamerata</i>	
4.6	4	13.5	10	3.5	4.3	12	16.6	16	18	17.2	17.6	19	19	7	5	4	<i>Pseudoguembelina costulata.</i>	
0	0	0	0	0	0	0	0	0	0	0	0	0	0	4.5	2.2	8	<i>Pseudoguembelina hariaensis</i>	
3	1.8	1.5	2	2.5	2	8	5.4	10	11	9.8	6.4	8	3	7.5	5.3	4	<i>Pseudoguembelina palpebra</i>	
1.3	1.8	4	2.6	1.2	2.8	3.8	6.9	6	9.4	4.8	6.7	8	3.5	1.8	1	2.1	<i>Pseudotextularia elegans</i>	
0	0	0	0	0	0	0.3	12	14	8	11	14	16	12	8.3	11	10	<i>Racemiguembelina fructicosa</i>	
5.5	2.1	16	14	3.9	0	29	9.7	14	13	12	13	14	16	11	8.9	9.6	<i>Rugoglobigerina hexacamerata</i>	
0.6	0	6.9	7.5	5.1	0	3.4	4.2	4.3	4.8	5.5	2.1	2.8	5.1	5.3	5.6	5.5	<i>Rugoglobigerina macrocephala</i>	
9.7	8.8	20	17	9	18	23	9	5.1	6.7	8.2	7.9	6.4	13	16	22	19	<i>Rugoglobigerina rugosa</i>	
1.9	4.5	15	19	12	35	0.6	0.8	0.9	1.2	1.6	1.3	1	1.5	2	0.4	1.6	<i>Rugoglobigerina rotundata</i>	

Globotruncanita falsocalcarata, *G. stuarti*, *G. stuartiformis*, *Heterohelix globulosa*, *Planoglobulina acervulinoides*, *P. multicamerata*, *Pseudoguembelina palpebra*, *P. costulata*, *P. palpebra*, *Pseudotextularia elegans*, *Racemiguembelina fructicosa*, *Rugoglobigerina hexacamerata*, *R. macrocephala*, *R. rugosa*, *R. rotundata*.

This zone has been identified in the Akveren Formation in samples K48–K50 of the Karainler section (Table 1, Figure 3), as well as in sample KD1 from the lowermost levels of the Karadur section (Table 2, Figure 5).

Plummerita hantkeninoides Zone is absent from all the Bartın sites which evolved within the last 300 k.y. of the Maastrichtian (Abramovich and Keller, 2002). The markers species of this zone is apparently preferred to live in eutrophic shelfal to upper-slope continental margin environments (e.g., Abramovich et al. 1998; Abramovich and Keller, 2002; MacLeod et al., 2007) and has not been reported from open-ocean pelagic carbonate sediments (Huber et al., 2008).

The Cretaceous-Paleogene transition is within the Akveren Formation. Any lithological changes were not seen in this region. Consequently, this boundary was estimated with changes of planktonic foraminifera species of this area.

K/Pg boundary are defined planktonic foraminiferal zone *Guembelitra cretacea* (P0) followed by the *Parvularugoglobigerina eugubina* Zone (P1a) (Smit, 1982). Abramovich et al. (2010) studied the distribution of the *Guembelitra* genera in detail in the uppermost Maastrichtian. They mentioned that whereas *Guembelitra* genera bloomed in the uppermost surface water primarily above shelf and slope environments, it appears to have failed to reach the open ocean.

5.2.2 Paleocene Planktonic Foraminifera Zones

Parvularugoglobigerina eugubina Taxon-range Zone (Pα): Biostratigraphic interval characterized by the total range of the nominate taxon *Parvularugoglobigerina eugubina*.

Author: Luterbacher and Premoli Silva, 1964.

Age: 64.97–64.8 Ma; Danian (early Paleocene).

The most characteristic assemblage recorded in this zone includes *Eoglobigerina eobulloides*, *Globoconusa conusa* and *Parvularugoglobigerina eugubina*.

This zone has been identified in sample KD2 of the Karadur section of the Akveren Formation (Table 2, Figure 5).

Parasubbotina pseudobulloides Partial-range Subzone (P1a) and *Subbotina triloculinoides* Lowest Occurrence Subzone (P1b) were not determined in this study area (Table 2, Figure 5).

Glomobanomalina compressa/*Praemurica uncinata* Lowest Occurrence Subzone (P1c): Biostratigraphic interval between the FO of *Glomobanomalina compressa* and/or *Praemurica inconstans* and the FO of *Praemurica uncinata*.

Author: Berggren and Miller, 1988.

Age: 62.87–61.37 Ma; early Paleocene (mid-late Danian).

The characteristic assemblage recorded in this zone includes *Globoconusa conusa*, *Parasubbotina pseudobulloides*, *Eoglobigerina eobulloides*, *Subbotina triloculinoides*, *Praemurica inconstans*, *Morozovella trinidadensis* and *Glomobanomalina compressa*.

This zone has been identified in sample KD3-KD11 of the Karadur section (Table 2, Figure 5).

Praemurica uncinata Lowest Occurrence Zone (P2): Biostratigraphic interval between the FO of *Praemurica uncinata* and the FO of *Morozovella angulata*.

Author: Berggren and Miller, 1988.

Age: 61.37–61.0 Ma; late early Paleocene (late Danian).

Associated species of the nominate taxon are *Globoconusa conusa*, *Parasubbotina pseudobulloides*, *Subbotina triloculinoides*, *Praemurica inconstans*, *Morozovella trinidadensis*, *Glomobanomalina compressa* and *Praemurica uncinata*.

This zone has been identified in sample KD12-

Table 2- Percentage abundances (%) of planktonic foraminifera are represented against sample position of the Karadur stratigraphic section from Maastrichtian to Selandian.

Maastr.	Danian										Selandian					Stage						
	<i>G. compressa</i>										<i>M. angulata</i>					Planktonic Foraminifera Zones						
	<i>P. panchinata</i>										<i>P. eugubina</i>											
Pharicensis	KD1										KD2					Samples						
0.5	2.1	0.6	0.3	0.5	2.2	0.7	1.3	0.6	0.3	4.2	0.2	0.7	4.5	3.2	1.0	3.4	1.1	4.7	2.6	4.2	4.7	<i>Abathomphalus mayaroensis</i>
																						<i>Contusotruncana contusa</i>
																						<i>Contusotruncana plummerea</i>
																						<i>Contusotruncana walfischensis</i>
																						<i>Gansserina gansseri</i>
																						<i>Globigerinelloides ultramicrus</i>
																						<i>Globotruncana aegyptiaca</i>
																						<i>Globotruncana arca</i>
																						<i>Globotruncana falsostuarti</i>
																						<i>Globotruncanella havanensis</i>
																						<i>Globotruncanella petaloidea</i>
																						<i>Globotruncanita stuarti</i>
																						<i>Globotruncanita stuartiformis</i>
																						<i>Heterohelix globulosa</i>
																						<i>Planoglobulina acervulinoides</i>
																						<i>Pseudoguembelina costulata.</i>
																						<i>Pseudoguembelina hariaensis</i>
																						<i>Pseudoguembelina palpebra</i>
																						<i>Racemiguembelina fructicosa</i>
																						<i>Rugoglobigerina hexacamerata</i>
																						<i>Rugoglobigerina macrocephala</i>
																						<i>Rugoglobigerina rugosa</i>
																						<i>Parvularugoglobigerina eugubina</i>
																						<i>Eoglobigerina eobulloides</i>
																						<i>Globoconusa conusa</i>
																						<i>Parasubbotina pseudobulloides</i>
																						<i>Subbotina triloculinoides</i>
																						<i>Glomobanomalina compressa</i>
																						<i>Praemucira inconstans</i>
																						<i>Morozovella trinidadensis</i>
																						<i>Praemurica uncinata</i>
																						<i>Morozovella angulata</i>
																						<i>Morozovella conicotruncana</i>
																						<i>Igorina pusilla</i>
																						<i>Morozovella velascoensis</i>
																						<i>Acarinina primitiva</i>
																						<i>Planorotalites chapmani</i>

KD13 of the Karadur section (Table 2, Figure 5) of the Akveren Formation.

Morozovella angulata Lowest Occurrence Zone (P3): Biostratigraphic interval between the FO of *Morozovella angulata* and the FO of *Globanomalina pseudomenardii*.

Author: Berggren and Miller, 1988.

Age: 61.0–59.4 Ma; early late Paleocene (Selandian).

Associated species of the nominate taxon in the studied section are *Globoconusa conusa*, *Parasubbotina pseudobulloides*, *Subbotina triloculinoides*, *Praemurica inconstans*, *Morozovella trinidadensis*, *M. angulata*, *M. conicotruncana*, *M. velascoensis*, *Glomobanomalina compressa*, *Praemurica uncinata*, *Acarinina primitiva*, *Igorina pusilla* and *Planorotalites chapmani*.

This zone has been identified samples KD14–KD20 of the Akveren Formation (Table 2, Figure 5).

5.3. Planktonic Foraminifera Assemblages

5.3.1. Composition of the Planktonic Foraminifera Assemblages in the Maastrichtian

The Maastrichtian is represented by small-sized biserial and low trochospiral spired planktonic foraminifera in the study area. The most dominant species in the Maastrichtian is small-sized *Rugoglobigerina* (*Rugoglobigerina rugosa*, *R. hexacamerata*, *R. macrocephala*) with low trochospiral shell. Within the foraminifera assemblages, *Rugoglobigerina* spp. is the most dominant species with 10-55% enrichment (Table 1, Figure 6). *Rugoglobigerina rotundata* with

higher trochospiral shell is the only dominant species in the lower Maastrichtian with abundance of 45% in sample K33 and 35% in sample K38. Although it has abundance between 2% and 45% in the lower Maastrichtian, the species is rare in the upper Maastrichtian (0.4-2%). *Pseudoguembelina* spp. is the second most dominant species (6-27%) (Table 1, Figure 6).

The third dominant species is *Heterohelix globulosa* with abundance of 2-16%. The other small-sized and low trochospiral shell species *Globotruncanella petaloidea* is another dominant species (2-17%). *Pseudotextularia elegans* is observed as a less common species (1-7%). In the upper Maastrichtian, *Racemiguembelina fructicosa* is the second common species (8-14%) together with *Rugoglobigerina* spp., with the exception of sample K40 (Table 1, Figure 6).

5.3.2. Composition of the Planktonic Foraminifera Assemblages in the Paleocene

Planktonic foraminifera species appearing just after the K/Pg boundary are small sized with globular chambers. The first Danian genera are *Parvularugoglobigerina eugubina*, *Eoglobigerina eobulloides* and *Subbotina triloculinoides*. In the lower Danian, *Globoconusa conusa*, *Parasubbotina pseudobulloides*, *Glomobanomalina compressa* and *Praemurica inconstans* are also observed together with *P. eugubina*, *E. eobulloides* and *S. triloculinoides* (Table 2, Figure 7).

Beginning from upper Danian, *Morozovella* spp. species appear (Table 2, Figure 7). The life time of *P. eugubina* and *E. eobulloides*, which are the initial species appearing in the Danian, is short. They are observed only in a few samples (Figure 7). The most dominant species of the Danian is *S. triloculinoides*

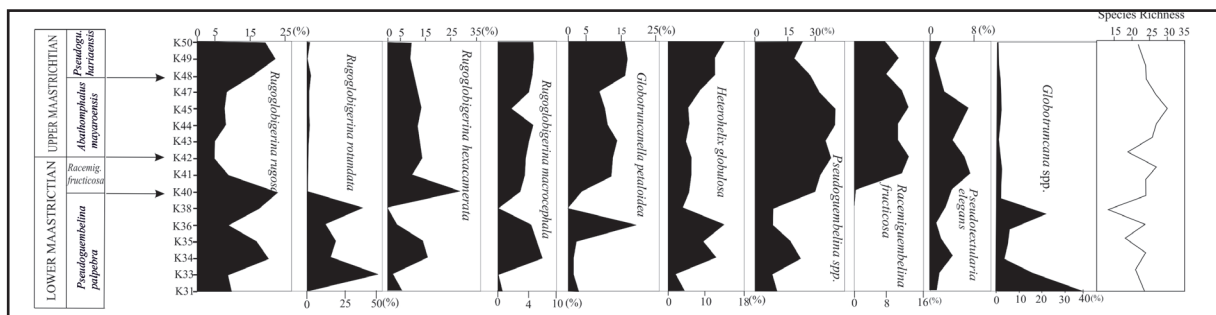


Figure 6- Species richness, percentage distributions of selected planktonic foraminifera of the Karainler stratigraphic section in the Maastrichtian.

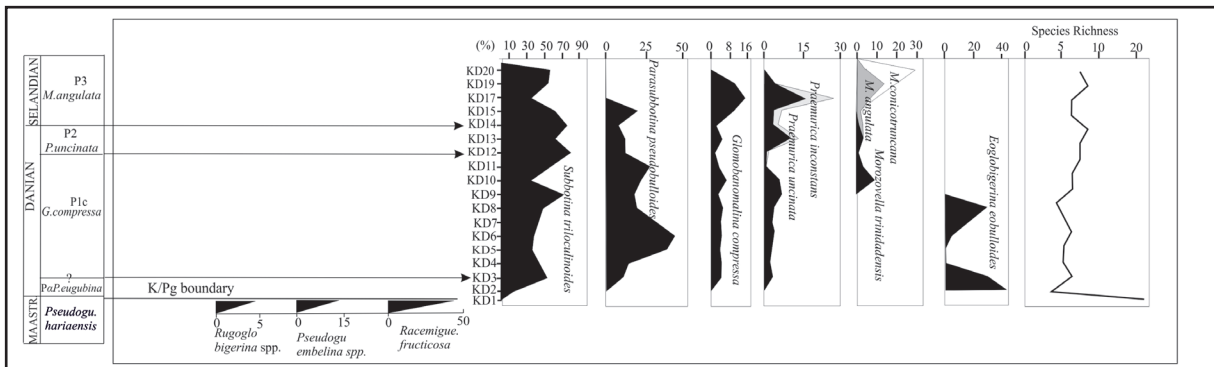


Figure 7- Species richness, percentage distributions of selected planktonic foraminifera of the Karadur stratigraphic section from Maastrichtian to Selandian.

with 14-79% abundance. *P. pseudobulloides* is the other dominant species with 8-43% abundance (Table 2, Figure 7).

Less dominant species are *G. compressa* (2-14%), *P. inconstans* (3-21%), *G. conusa* (2-7%), *Morozovella trinidadensis* (1-9%), *Praemurica uncinata* (1-14%), *Morozovella angulata* (2-7%) and *Morozovella conicotruncana* (10-15%). In the Selandian, *S. triloculinoides* (53-57%) and *Morozovella* species are dominant (Table 2, Figure 7).

5.4. The Paleocology of Planktonic Foraminifera in the Maastrichtian

Planktonic foraminifera are defined as sensitive recorders of the upper part of the water column, reflecting changes in primary productivity, sea surface water temperature and the degree of water column stratification (Leckie, 1987; Schiebel and Hemleben, 2000). In the Bartın area, abundance of *Globotruncana* spp. (*Globotruncana arca*, *G. linneiana*, *G. aegyptiaca*, *G. mariei*, *G. bulloides*, *G. falsostuarti*, *G. ventricosa* and *G. insignis*) is rare and shows a decrease in the Maastrichtian (Table 1, Figure 6). Abramovich and Keller (2003) indicated that the abundance of globotruncanids decreased under biotic stress in the Maastrichtian. In this study area, decrease in the abundance of globotruncanids may reflect the changes in environmental conditions.

In the lower Maastrichtian, the abundance of *Rugoglobigerina* taxa increased when abundances of globotruncanids decreased. Genus *Rugoglobigerina* are widely observed from lower Maastrichtian to upper Maastrichtian (Table 1, Figure 6). This genus was proposed to live in tropical-mid latitudes and

warm climates in previous studies and the abundance of *Rugoglobigerinids* arrive at maximum levels in climatic warming periods (Malmgren, 1991; Nederbragt, 1991, 1998; Li and Keller, 1998a). Abramovich and Keller (2003) stated that decreases in the relative abundance of tropical-subtropical *Pseudoguembelina costulata*, *Rugoglobigerina rugosa* and several keeled globotruncanids (including *Contusotruncana* species) coincide with the warm event and their decline suggests unfavorable environmental conditions.

Pseudoguembelina and *Heterohelix globulosa* are dominant in the upper Maastrichtian fauna in the study area (Table 1, figure 6). Abramovich et al. (2003) and Keller (2004) suggest that the domination of small biserial ecological species and low trochospiral species (*Rugoglobigerina*, *H. globulosa*, *Pseudoguembelina*, *Pseudotextularia*) and also the low diversity of the species indicates the presence of highly stressed environmental conditions.

Heterohelix globulosa is evaluated as tolerant of the changes in salinity, temperature, oxygen and the amount of nutrition (Keller, 2004). *Heterohelix globulosa* continued to live without being affected by changes in environmental conditions. Ashckenazi-Polivoda et al. (2011) stated that Heterohelicids display high abundance in a variety of marine environments, indicate unsuitable environmental conditions and an ability to adjust to a wide range of water column conditions. In this study, increasing abundance of *Heterohelix globulosa* and *Pseudoguembelina* spp. shows that they are resistant and tolerant of the changes in environmental conditions such as nutrition and temperature, exhibiting compatibility with global climate changes before the K/Pg boundary.

Racemiguembelina, which is common in the upper Maastrichtian assemblages, were defined as photosymbiotic (Houston and Huber, 1998; D'Hondt and Zachos, 1998; Houston et al., 1999). *Racemiguembelina* is defined as an oligotrophic genus in the open ocean (Abramovich et al., 2010). In the uppermost Maastrichtian, high abundance of *Racemiguembelina fructicosa*, *Heterohelix globulosa* and *Rugoglobigerina* spp. can be interpreted as the presence of oligotrophic and warmer environmental conditions in this study (Table 1-2, Figure 6-7).

Kaya-Özer (2014) investigated nanofossil assemblages, carbon and oxygen stable isotope data in this stratigraphic section and suggested that environmental conditions changed from eutrophic to oligotrophic during the Maastrichtian. Diversification of planktonic foraminifera species indicate the same environmental conditions in this study.

In the upper Maastrichtian in the Bartın area, the increasing abundance of *Globotruncanella petaloidea* is coherent with increasing abundance of the species which are tolerant of changes in nutrition and temperature (Table 1, Figure 6). It is possible that these species are able to live in similar environments.

5.5. The Paleocology of Planktonic Foraminifera in the Paleocene

Small, unornamental and primitive Paleogene species began to be seen just above K/Pg boundary in this study area. *Parasubbotina pseudobulloides* and *Subbotina triloculinoidea* are the dominant species in the Danian (Table 2, Figure 7). In the upper Danian, together with these species, the abundance of *Praemurica* species also increased. However, in the Selandian, *Morozovella* species start to dominate (Table 2, Figure 7).

Quillevère and Norris (2003) suggested that different depth habitats played an important role in the diversification of planktonic foraminifera, related to the initiation of photosymbiosis as a trophic strategy. Subbotinids lived within or below the thermocline whereas morozovellids and igorinids dwelled in the surface mixed-layer (Shackleton et al., 1985; Pearson et al., 1993; D'Hondt et al., 1994; Van Eijden, 1995; Lu et al., 1998; Quillevère and Norris, 2003).

Subbotina and *Parasubbotina* preferred cold water, asymbiotic living and mesotrophic environmental

conditions (Boersma and Premoli-Silva, 1991; Pearson et al., 1993; Norris, 1996; Berggren and Norris, 1997; Guasti et al., 2005). Also, *Parasubbotina* and *Subbotina* taxa shared a similar trophic strategy and living position within the thermocline (Guasti et al., 2005). Guasti et al. (2005) stated that increased productivity through enhanced input of nutrients from land may favor the thermocline dweller *Parasubbotina*. In the study area, during the Danian, high abundance of *Parasubbotina* and *Subbotina* can be considered as indicating mesotrophic and cooler environmental conditions.

Stable isotope values indicate that subbotinids lived in cooler, deeper waters than morozovellids (Boersma and Premoli Silva, 1991; Norris, 1996). Positive $\delta^{18}\text{O}$ values show *Morozovella* and later *Acarinina* species have a deeper habitat (Berggren and Norris, 1997; Quillevère et al., 2000; Quillevère and Norris, 2003). Most morozovellids, acarininids and igorinids were accepted as photosymbiotic species (D'Hondt et al., 1994; Norris, 1996; Berggren and Norris, 1997; Quillevère and Norris, 2003). Acarininids and morozovellids thrive in low-nutrient water masses in the global ocean during the late Paleocene (Norris, 1996; Quillevère and Norris, 2003). *Praemurica inconstans* was accepted as a photosymbiotic species (Kelly et al., 1996).

Morozovella adapted to warm water and oligotrophic environmental conditions (Pearson et al., 1993; Norris, 1996; Berggren and Norris, 1997; Quillevère and Norris, 2003). In this study, in the Selandian, *Morozovella* started with common taxa with *Subbotina* and *Parasubbotina* (Table 2, Figure 7). Increasing abundance of *Morozovella* taxa which lived in low nutrition waters, indicates mesotrophic-oligotrophic and warmer environmental conditions in the Selandian. Kaya-Özer (2014) suggested similar environmental conditions in the Paleocene based on $\delta^{13}\text{C}$ values.

6. Conclusion

The Akveren Formation represents open sea characteristics in the Maastrichtian-Selandian period based on planktonic foraminifera species. In the Maastrichtian, planktonic foraminifera assemblages in the study area are dominated by small, simple morphotypes, opportunists, generalists and high-stress specialist morphologies. Increased abundance

of small-sized, biserial and low trochospiral planktonic foraminifera species (*Rugoglobigerina* spp., *Heterohelix globulosa*, *Pseudoguembelina* spp., *Pseudotextularia elegans*) show variable and unstable environmental conditions in the Western Black Sea basin during the Maastrichtian. In the uppermost Maastrichtian, high abundance of *Racemiguembelina fructifera*, *Heterohelix globulosa* and *Rugoglobigerina* spp. indicates oligotrophic and warmer environmental conditions.

In the lower Paleocene, high abundance of *Parasubbotina* and *Subbotina* shows mesotrophic and cooler environmental conditions. In the Selandian, increasing abundance of *Morozovella* taxa indicates mesotrophic- oligotrophic and warmer environmental conditions.

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References

- Abramovich, S., Almogi-Labin, A., Benjamini, C. 1998. Decline of the Maastrichtian pelagic ecosystem based on planktic foraminifera assemblage changes: Implication for the terminal Cretaceous faunal Crisis. *Geology* 26, 63-66.
- Abramovich, S., Keller, G. 2002. High stress late Maastrichtian paleoenvironment: Inference from planktonic foraminifera in Tunisia, *Palaeogeography, Palaeoclimatology, Palaeoecology* 178, 145-164.
- Abramovich, S., Keller, G. 2003. Planktonic foraminiferal response to the latest Maastrichtian abrupt warm event: a case study from South Atlantic DSDP Site 525A. *Marine Micropaleontology* 48, 225-249.
- Abramovich, S., Keller, G., Stüben, D., Berner, Z. 2003. Characterization of late Campanian and Maastrichtian planktic foraminiferal depth habitats and vital activities based on stable isotopes. *Palaeogeography, Palaeoclimatology, Palaeoecology* 202, 1-29.
- Abramovich, S., Yovel-Corem, S., Almogi-Labin, A., Benjamini C. 2010. Global climate change and planktic foraminiferal response in the Maastrichtian. *Paleoceanography* 25, 1-15.
- Akman, Ü. 1992. Amasra-Arıt arasının jeolojisi. Doktora Tezi, Ankara Üniversitesi, 209 s. Ankara, (unpublished).
- Akyol, Z., Arpat, E., Erdoğan, B., Göğler, E., Güner, Y., Şaroğlu, F., Şentürk, İ., Tütüncü, K., Uysal, Ş. 1974. 1/50.000 ölçekli Türkiye Jeoloji haritası serisi, Zonguldak E29a, E29b, E29c, E29d, Kastamonu E30a, E30d. Maden Tetkik ve Arama Genel Müdürlüğü Yayınları, Ankara.
- Ashckenazi-Polivoda, S., Abramovich, S., Almogi-Labin, A., Schneider-Mor, A., Feinstein, S., Puttmann, W., Berner A. 2011. Paleoenvironments of the latest Cretaceous oil shale sequence, Southern Tethys, Israel, as an integral part of the prevailing upwelling system. *Palaeogeography, Palaeoclimatology, Palaeoecology* 305, 93-108.
- Aydın, A. 2005. İzmit kuzeybatısı Geç Kretase-Paleojen nannoplankton biyostratigrafisi. Yüksek lisans tezi, Ankara Üniversitesi, 237 s. Ankara, (unpublished).
- Aydın, M., Şahintürk, Ö., Serdar, H.S., Özçelik, Y., Akarsu, İ., Üngör, A., Çokuğraş, R., Kasar, S. 1986. Ballıdağ-Çangaldağ (Kastamonu) arasındaki bölgenin jeolojisi. *Türkiye Jeoloji Kurumu Bülteni* 29, 1-16.
- Badgley, P.C. 1959. Sinop Havzasının petrol olanakları. Petrol işleri Genel Müdürlüğü arşivi, Ankara.
- Barka, A., Sütçü, Y. F., Tekin, F., Gedik, İ., Karabıyıkoglu, M., Saraç, G. Önal, Ö., Arel, E., Özdemir M. 1983. Sinop Yarımadasının jeolojisi ve tektonik evrimi. *Türkiye Jeoloji Kurumu Bülteni* 4, 24.
- Barrera, E., Savin, S.M., Thomas, E., Jones, C.E. 1997. Evidence for thermohaline circulation reversals controlled by sea-level change in the latest Cretaceous. *Geology* 25, 715-718.
- Barrera, E., Savin, S.M. 1999. Evolution of late Campanian-Maastrichtian marine climates and oceans. Barrera, E., Johnson, C.C. (Ed.). *Evolution of the Cretaceous Ocean-Climate System. Geological Society of America Special Paper* 332, 245-282.
- Berggren, W.A., Norris, R.D. 1997. Biostratigraphy, phylogeny and systematics of Paleocene trochospiral planktic foraminifera. *Micropaleontology* 43, 1-116.

- Berggren, W.A., Miller, K.G. 1988. Paleocene tropical planktonic foraminiferal biostratigraphy and magneto biochronology. *Micropaleontology* 34, 4, 362-380.
- Bice, K.L., Birgel, D., Meyers, P.A., Dahl, K., Hinrichs K, U., Norris, R.D. 2006. A multiple proxy and model study of Cretaceous upper ocean temperatures and atmospheric CO2 concentrations. *Paleoceanography* 21, 1-17.
- Boersma, A., Premoli Silva, I. 1991. Distribution of Paleogene planktonic foraminifera analogies with Recent? *Palaeogeography, Palaeoclimatology, Palaeoecology* 83, 29-48.
- Bornemann, A., Norris, R.D., Friedrich, O., Beckmann, B., Schouten, S., Sinninghe Damste, J.S., Vogel, J., Hofmann, P., Wagner, T. 2008. Isotopic evidence for glaciation during the Cretaceous supergreen house. *Science* 319, 189-192.
- Caron, M. 1985. Cretaceous planktic foraminifera. Bolli H.M., Saunders J.B., Perch-Nielsen, K. (Ed.). *Plankton Stratigraphy*. Cambridge University Press, London, 1032 p.
- Clarke, L.J., Jenkyns, H.C. 1999. New oxygen isotope evidence for long-term Cretaceous climatic change in the Southern Hemisphere. *Geology* 27, 8, 699-702.
- Corfield, R.M., Norris, R.D. 1996. Deep water circulation in the Paleocene ocean. Knox R., Corfield, R.M., Dunay, R.E. (Ed.). *Correlation of the early Paleogene in Northwest Europe*. Geological Society of London Special Publication 101, 443-456.
- Deconto, R.M., Hay, W.W., Thompson, S.L., Berggren, J. 1999. Late Cretaceous climate and vegetation interactions: Cold continental interior paradox. Barrera, E., Johnson, C.C. (Ed). *Evolution of the Cretaceous Ocean-Climate System*. Boulder, CO: Geological Society of America Special Paper 332, 391-406.
- Derman, A. S. 2002. Karadeniz'in açılma istifleri (Black Sea rift sequences). *Turkish Association of Petroleum Geologists Bulletin* 14, 37-66.
- Derman, A.S., Alişan, C., Özçelik, Y. 1995. Himmetpaşa Formation: new palynological age data and its significance. Erler, A., Ercan, T., Bingöl, E., Örçen, S. (Eds). *Geology of the Black Sea region*. General Directorate of Mineral Research and Exploration and Chamber of Geological Engineers 99-103.
- D'Hondt, S., Lindinger, M. 1994. A stable isotopic record of the Maastrichtian oceanclimate system: South Atlantic DSDP Site 528. *Palaeogeography, Palaeoclimatology, Palaeoecology* 112, 363-378.
- D'Hondt, S., Zachos, J.C., Schultz, G. 1994. Stable Isotopic signal and Photosymbiosis in late Paleocene Planktic foraminifera. *Paleobiology* 30, 3, 391-406.
- D'Hondt, S., Zachos, J. 1998. Cretaceous foraminifera and the evolutionary history of planktic photosymbiosis. *Paleobiology* 24, 512-523.
- Dizer, A., Meriç, E. 1981. Kuzeybatı Anadolu'da Üst Kretase-Paleosen biyostratigrafisi. *Maden Tetkik ve Arama Dergisi* 95-96, 149-163.
- Dobrucalı, S. 1985. Akveren Formasyonu (Sinop-Gerze) planktonik foraminifer faunası ve stratigrafik konumu. Yüksek lisans tezi, Karadeniz Teknik Üniversitesi, 56, Trabzon, (unpublished).
- Forster, A., Schouten, S., Moriya, K., Wilson, P.A., Sinninghe Damste, J.S. 2007. Tropical warming and intermittent cooling during the Cenomanian/Turonian oceanic anoxic event 2: Sea surface temperature records from the equatorial Atlantic. *Paleoceanography* 22, 1-14.
- Friedrich, O., Erbacher, J., Moriya, K., Wilson, P.A., Kuhnert, H. 2008. Warm saline intermediate waters in the Cretaceous tropical Atlantic Ocean. *Nature Geoscience* 1, 453-457.
- Friedrich, O., Norris, R.D., Erbacher, J. 2012. Evolution of middle to Late Cretaceous oceans-A 55 m.y. record of Earth's temperature and carbon cycle. *Geology* 40, 107-110.
- Gayle, R.B. 1959. Sinop Yöresi ile ilgili çalışma. *Petrol işleri, Rapor no.17*, Ankara.
- Gedik, A., Ercan, T., Korkmaz, S. 1983. Orta Karadeniz (Samsun-Sinop) havzası jeolojisi ve volkanik kayaların petrolojisi. *Maden Tetkik ve Arama Dergisi* 99-100, 34-51.
- Gedik, A., Korkmaz, S. 1984. Sinop Havzasının jeolojisi ve petrol olanakları. *Jeoloji Mühendisleri Odası Dergisi* 19, 53-79.
- Gedik, I., Timur, E., Duru, M., Pehlivan, S. 2005. 1:50.000 Ölçekli Türkiye Jeoloji Haritaları (İstanbul F22 c, d; F23 c, d; Bursa G22 a, b; G23 a, b), Maden Tetkik ve Arama Genel Müdürlüğü Jeoloji Etütleri Dairesi, No: 10-17, Ankara.

- Görür, N. 1988. Timing of opening of the Black Sea basin. *Tectonophysics* 147, 247-262.
- Görür, N., Tüysüz, O. 1997. Petroleum geology of the southern continental margin of the Black Sea. Robinson, A. G. (Ed). *Regional and petroleum geology of the Black Sea and surrounding region*. American Association of Petroleum Geologists (AAPG) Memoirs 68, 227-240.
- Guasti, E., Speijer, R. P., Fornaciari, E., Schmitz, B., Kroon, D., Gharaibeh, A. 2005. Transient biotic change within the Danian-Selandian transition in Egypt and Jordan. In: *Early Paleogene environmental turnover in the Southern Tethys as recorded by foraminiferal and organic-walled dinoflagellate cysts assemblages*. *Berichte aus dem Fachbereich Geowissenschaften der Universität Bremen*, 241, 75-110.
- Güray, A. 2006. Campanian-Maastrichtian planktonic foraminiferal investigation and biostratigraphy (Kokaksu section, Bartın, NW Anatolia): remarks on the Cretaceous paleoceanography based on quantitative data. Yüksek lisans tezi, Orta Doğu Teknik Üniversitesi, 244, Ankara, (unpublished).
- Houston, R.M., Huber, B.T. 1998. Evidence of photosymbiosis in fossil taxa? Ontogenetic stable isotope trends in some late Cretaceous planktonic foraminifera. *Marine Micropaleontology* 34, 29-46.
- Houston, R.M., Huber, B.T., Spero, H.J. 1999. Size-related isotopic trends in some Maastrichtian planktonic foraminifera: methodological comparisons, intraspecific variability and evidence for photosymbiosis. *Marine Micropaleontology* 36, 169-188.
- Huber, B.T., Hodell, D.A., Hamilton, C.P. 1995. Middle-Late Cretaceous climate of the southern high latitudes: Stable isotopic evidence for minimal equator-to-pole thermal gradients. *GSA Bulletin* 107, 1164-1191.
- Huber, B.T., Macleod, K.G., Norris, R.D. 2002. Abrupt Extinction and Subsequent Reworking of Cretaceous Planktonic Foraminifera across the Cretaceous-Tertiary Boundary: Evidence from the Subtropical North Atlantic. *Geological Society of America* 356, 277-289.
- Huber, B.T., Macleod, K.G., Tur, N.A. 2008. Chronostratigraphic framework for Upper Campanian-Maastrichtian sediments on the Blake Nose (Subtropical North Atlantic). *Journal of Foraminiferal Research* 38, 162-182.
- Kaya, O., Dizer, A., Tansel, İ., Meriç, E. 1982/1983. Ereğli (Zonguldak) alanının Kretase stratigrafisi. *Maden Tetkik ve Arama Dergisi* 99/100, 19-33.
- Kaya-Özer, C. 2009. Bartın-Kurucaşile yöresi Geç Kretase birimleri planktonik foraminifer ve nannoplankton biyostratigrafisi. Doktora Tezi, Ankara Üniversitesi, Fen Bilimleri Enstitüsü, 371, Ankara (unpublished).
- Kaya-Özer, C. 2014. Calcareous Nannofossil Assemblages Changes and Stable Isotope Data from Maastrichtian to Selandian in the Akveren Formation, Western Black Sea, Turkey. *Arabian Journal of Geosciences* 7, 1233-1247.
- Kaya-Özer, C., Toker, V. 2012. Calcareous Nannoplankton Biostratigraphy of the Bartın Province, Western Black Sea, Turkey. *Acta Geologica Sinica - English Edition* 86, 6, 1434-1446.
- Keller, G. 1996. The K/T mass extinction in planktic foraminifera biotic constraints for catastrophe theories. Mac-Leod, N., Keller, G., (Ed.) *The Cretaceous-Tertiary mass extinction: Biotic and environmental events*, Norton Press New York 63-100.
- Keller, G. 2004. Low-diversity, Late Maastrichtian and Early Danian planktic foraminiferal assemblages of the Eastern Tethys. *Journal of Foraminiferal Research* 34, 1, 49-73.
- Keller, G., Adatte, T., Stinnesbeck, W., Luciani, V., Karoui-Yaakoub, N., Zaghbib-Turki, D. 2002. Paleoecology of the Cretaceous-Tertiary mass extinction in planktonic foraminifera. *Palaeogeography, Palaeoclimatology, Palaeoecology* 178, 257-297.
- Kelly, C. D., Arnold, A. J., Parker, W. C. 1996. Paedomorphosis and the origin of the Paleogene planktonic foraminiferal genus *Morozovella*. *Paleobiology* 22, 2, 266-281.
- Ketin, İ., Gümüş, A. 1963. Sinop-Ayancık arasında III. Bölgeye dahil sahalarnın jeolojisi. TPAO, Rapor no. 288. Ankara.

- Kırcı, E. 1998. Cide (Kastamonu) yöresinde Kretase-Tersiyer geçişinin biyostratigrafisi. Yüksek lisans tezi, İstanbul Üniversitesi, 77, İstanbul, (unpublished).
- Leckie, R.M. 1987. Paleoecology of mid-Cretaceous planktonic foraminifera: A comparison of open ocean and epicontinental sea assemblages. *Micropaleontology* 33, 264-276.
- Li, L., Keller, G. 1998a. Maastrichtian climate productivity and faunal turnovers in planktic foraminifera in South Atlantic DSDP Sites 525A and 21. *Marine Micropaleontology* 33, 55-86.
- Li, L., Keller, G. 1998b. Diversification and extinction in Campanian-Maastrichtian planktic foraminifera of Northwestern Tunisia. *Ecl. Helv* 91, 75-102.
- Lu, G., Adatte, T., Keller, G., Ortiz, N. 1998. Abrupt climatic, oceanographic and ecologic changes near the Paleocene-Eocene transition in the deep Tethys basin: The Alamedilla section, southern Spain. *Eclogae geol. Helv.* 91, 293-306.
- Luterbacher, H.P., Premoli Silva, I. 1964. Biostratigrafi a del limite cretaceo-terziario nell' appennino centrale. *Riv Ital Paleontol Stratigr* 70,67-128.
- Macleod, K.G., Whitney, D.L., Huber, B.T., Koeberl, C. 2007. Impact and extinction in remarkably complete Cretaceous-Tertiary boundary sections from Demerara Rise, tropical western north Atlantic. *Geological Society of America, Bulletin* 119, 101-115.
- Malmgren, B.A. 1991. Biogeographic patterns in terminal Cretaceous planktonic foraminifera from Tethyan and warm transitional waters. *Marine Micropaleontology* 18, 73-99.
- Miller, K.G., Wright, J.D., Browning, J.V. 2005. Visions of ice sheets in a greenhouse world. *Marine Geology* 217, 215-231.
- Nederbragt, A.J. 1991. Late Cretaceous biostratigraphy and development of Heterohelicidae (planktic foraminifera). *Micropaleontology* 37, 329-372.
- Nederbragt, A.J. 1998. Quantitative biogeography of late Maastrichtian planktic foraminifera. *Micropaleontology* 44, 385-412.
- Norris, R.D. 1996. Symbiosis as an Evolutionary Innovation in the radiation of Paleocene planktic foraminifera. *Paleobiology* 22, 4, 461-480.
- Norris, R.D., Bice, K.L., Magno, E.A., Wilson, P.A. 2002. Jiggling the tropical thermostat in the Cretaceous hothouse. *Geology* 30, 299-302.
- Olsson, R.C., Hemleben, C., Berggren, W.A., Huber, B.T. 1999. Atlas of Paleocene planktonic foraminifera. *Smithsonian Contributions to Paleobiology* 85, 1-252
- Özçelik, Y., Çaptuğ, A. 1990. Amasra doğusu-Cide arasında kalan alanda yapılan saha gözlemleri ve revizyon çalışmaları. Türkiye Petrolleri Anonim Ortaklığı, Rapor No:2789, Ankara.
- Özkan-Altın, S., Özcan, E. 1999. Upper Cretaceous planktonic foraminiferal biostratigraphy from NW Turkey: calibration of the stratigraphic ranges of larger benthonic foraminifera. *Geological Journal* 34, 287-301.
- Pearson, P.N., Shackleton, N.J., Hall, M.A. 1993. Stable isotope paleoecology of middle Eocene planktonic foraminifera and multi-species isotope stratigraphy, DSDP Site 523, South Atlantic. *Journal of Foraminiferal Research* 23, 2, 123-140.
- Petrizzo, M.R., Falzoni, F., Premoli Silva, I. 2011. Identification of the base of the lower-to-middle Campanian Globotruncana ventricosa zone: comments on reliability and global correlations. *Cretaceous Research* 32, 387-405.
- Quillevère, F., Norris, R.D., Berggren, W.A., Aubry, M.P. 2000. 59.2 Ma and 56.5 Ma: two significant moments in the evolution of acariniids (planktonic foraminifera). *GFF Early Paleogene warm climates and biosphere Dynamics* 122, 131-132.
- Quillevère, F., Norris, R.D. 2003. Ecological development of acariniids (planktonic foraminifera) and hydrographic evolution of Paleocene surface waters. S.L. Wing et al. (Ed.) *Causes and consequences of globally warm climates in the early Paleogene*. The Geological Society of America, Boulder, Colorado 369, 223-238.
- Robaszynski, F., Caron, M., Gonzales-Donoso, J.M., Wonders, A.A.H. 1984 the European Working Group on Planktonic Foraminifera. Atlas of Late Cretaceous globotruncanids. *Revue Micropaleontology* 26, 145-305.

- Robaszynski, F., Caron, M. 1995. Foraminifères planctoniques du Cretace; commentaire de la zonation Europe-Mediterranee Bulletin de la Societe Geologique de France 166, 681-692.
- Robinson, A.G., Rudat, J. H., Banks, C. J., Wiles, R. L.F. 1996. Petroleum geology of the Black Sea. Marine and Petroleum Geology 13, 195-223.
- Saner, S. 1980. Batı Pontidlerin ve komşu havzaların oluşumlarının levha tektoniği kuramıyla açıklanması, Kuzeybatı Türkiye. Maden Tetkik ve Arama Dergisi 93-94, 1-19.
- Sarıca, N. 1993. Gökçeada (Kastamonu) yöresinde Kretase/Tersiyer sınırının planktik foraminiferlerle biyostratigrafik incelemesi. Türkiye Jeoloji Kurumu Bülteni 8, 329-345.
- Schiebel, R., Hemleben, C. 2000. Interannual variability of planktic foraminiferal populations and test flux in the eastern North Atlantic Ocean (JGOFS). Deep Sea Research Part II: Topical Studies in Oceanography 47, 9-11, 1809-1852.
- Shackleton, N.J., Corfield, R.M., Hall, M.A. 1985. Stable isotope data and the ontogeny of Paleocene planktonic foraminifera. Journal of Foraminiferal Research 15, 4, 321-336.
- Shirazi, M.P.N., Shams, P., Bahrami, M. 2013. Biostratigraphy and paleoecology of Maastrichtian sediments in the Zagros Basin, Iran. Acta Geologica Sinica (English Edition) 87, 5, 1387-1395.
- Sirel, E. 1991. Cide Bölgesi (Kuzey Turkey) Maastrichtiyende bulunan yeni bir foraminifer cinsi: Cideina n.gen. Maden Tetkik ve Arama Dergisi 112, 149-154.
- Smit, J. 1982. Extinction and evolution of planktonic foraminifera after a major impact at the Cretaceous/Tertiary boundary. Silver, L.T., Schultz, P.H. (Ed.) In Geological implications of impacts of large asteroids and comets on the Earth. Special Papers. Geological Society of America 190, 329-352.
- Sunal, G., Tüysüz, O. 2001. Batı Pontidler'de Tersiyer yaşlı çarpışma sonrası sıkışmalı yapıların özellikleri. Türkiye Petrol Jeologları Derneği Bülteni 13, 1-26.
- Şahintürk, O., Özçelik, Y. 1983. Zonguldak-Bartın-Amasra-Kurucaşile-Cide dolaylarının jeolojisi ve petrol olanakları. Türkiye Petrolleri Anonim Ortaklığı, Rapor No:1816, Ankara.
- Şener, S. 2007. Amasra yöresi (Batı Karadeniz) Üst Jura-Alt Kretase İnalıtı kireçtaşlarının bentik foraminifer paleontolojisi, Yüksek lisans tezi, Ankara Üniversitesi, 76, Ankara, (unpublished).
- Şengör, A.M.C. 1982. Ege'nin neotektonik evrimini yöneten etkenler, Batı Anadolu'nun genç tektoniği ve volkanizması paneli. Türkiye Jeoloji Kurultayı 59-71.
- Thibault, N., Gardin, S. 2007. The late Maastrichtian nannofossil record of climate change in the South Atlantic DSDP Hole 525A. Marine Micropaleontology 65, 163-184.
- Tokay, M. 1954/1955. Filyos çayı ağzı-Amasra-Bartın-Kozcağız-Çaycuma bölgesinin jeolojisi. Maden Tetkik ve Arama Dergisi 46/47, 58-74.
- Toumarkine, M., Luterbacher, H. 1985. Paleocene and Eocene planktic foraminifera. Bolli, H.M., Saunders, J.B., Perch-Nielsen, K. (Ed.) Plankton Stratigraphy. Cambridge University Press 87-154.
- Tüysüz, O. 1993. Karadeniz'den Orta Anadolu'ya bir Jeotravers: Kuzey Neo-Tetisin tektonik evrimi. Türkiye Petrol Jeologları Derneği Bülteni 5, 1-33.
- Tüysüz, O. 2002. Upper Cretaceous red pelagic limestones in the Pontides, northern Turkey and their significance on the geological evolution of Black Sea. Inaugural Workshop of IGCP 463, 30.
- Tüysüz, O., Kırıcı, S., Sunal, G. 1997. Cide-Kurucaşile dolayının jeolojisi. Türkiye Petrolleri Anonim Ortaklığı, Rapor No. 3736, Ankara.
- Tüysüz, O., Aksay, A., Yiğitbaş, E. 2004. Batı Karadeniz bölgesi litostratigrafi birimleri. Stratigrafi komitesi litostratigrafi birimleri serisi-1. Maden Tetkik ve Arama Genel Müdürlüğü Yayınları, Ankara.
- Tüysüz, O., Yılmaz, İ.Ö., Svabnicka, L., Kırıcı, S. 2012. The Unaz Formation: a key unit in the western Black Sea region, N Turkey. Turkish Journal of Earth Sciences 21, 1009-1028.
- Van Eijden, A.J.M. 1995. Morphology and relative frequency of planktic foraminiferal species in relation to oxygen isotopically inferred depth habitats. Palaeogeography, Palaeoclimatology, Palaeoecology 113, 267-301.

- Varol, O. 1983. The Cretaceous-Paleocene calcareous nannofossils from the Kokaksu section (Zonguldak, Northern Turkey). *Geol. Palaont. Abh.*166, 431-460.
- Wade, B.S., Pearson, P.N., Berggren, W.A., Palike, H. 2011. Review and revision of Cenozoic tropical planktonic foraminiferal biostratigraphy and calibration to the geomagnetic polarity and astronomical time scale. *Earth-Science Reviews* 104, 111–142.
- Wilson, P.A., Norris, R.D. 2001. Warm tropical ocean surface and global anoxia during the mid-Cretaceous period. *Nature* 412, 425-429.
- Wilson, P. A., Norris, R. D., Cooper, M. J. 2002. Testing the Cretaceous greenhouse hypothesis using glassy foraminiferal calcite from the core of the Turonian tropics on Demerara Rise. *Geology* 30, 607–610.
- Zachos, J.C., Arthur, M.A., Thunell, R.C., Williams, D. F., Tappa, E.T. 1985. Stable isotope and trace element geochemistry of carbonate sediments across the Cretaceous/Tertiary boundary at Deep Sea Drilling Project Hole 577. *Initial Reports of the Deep Sea Drilling Project* 86, 513-532.
- Zachos, J.C., Arthur, M.A., Dean, W.E. 1989. Geochemical evidence for suppression of pelagic marine productivity at the Cretaceous/Tertiary boundary. *Nature* 337, 61-64.

PLATES

PLATE I

Figure 1- *Abathomphalus mayaroensis* (Bolli), Spiral view, Sample K41.

Figure 2- *Abathomphalus mayaroensis* (Bolli), Umbilical view, Sample K41.

Figure 3- *Contusotruncana contusa* (Cushman), Spiral view, Sample K41.

Figure 4- *Gansserina gansseri* (Bolli), Side view, Sample K39.

Figure 5- *Rugoglobigerina rugosa* (Plummer), Spiral view, Sample K48.

Figure 6- *Heterohelix globulosa* (Ehrenberg), Side view, Sample K32.

Figure 7- *Planoglobulina multicamerata* (De Klasz), Side view, Sample K48.

Figure 8- *Pseudoguembelina hariaensis* Nederbragt, Side view, Sample K48.

Figure 9- *Pseudoguembelina hariaensis* Nederbragt, Side view, Sample K48.

Figure 10- *Pseudoguembelina palpebra* Bronnimann and Brown, Sample K36.

Figure 11- *Racemiguembelina fructicosa* (Egger), Side view, Sample K42.

Figure 12- *Pseudotextularia elegans* (Rzehak), Side view, Sample K48.

PLATE I

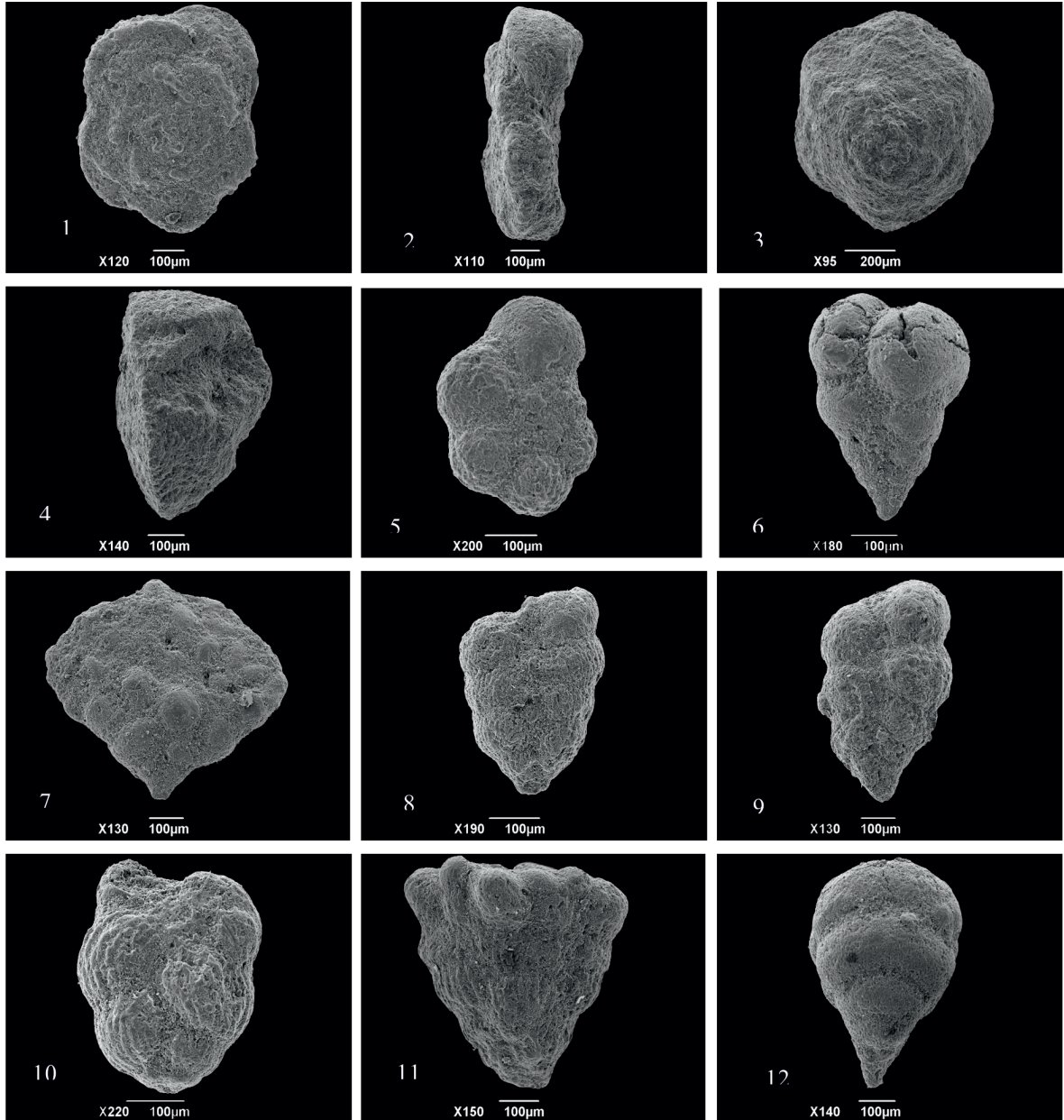


PLATE II

Figure 13- *Parvularugoglobigerina eugubina* (Luterbacher and Premoli Silva), Umbilical view, Sample KD3.

Figure 14- *Parasubbotina pseudobulloides* (Plummer), Spiral view, Sample KD6.

Figure 15- *Parasubbotina pseudobulloides* (Plummer), Side view, Sample KD15.

Figure 16- *Igorina pusilla* (Bolli), Spiral view, Sample KD20.

Figure 17- *Subbotina triloculinoides* (Plummer), Spiral view, Sample KD16.

Figure 18- *Subbotina triloculinoides* (Plummer), Umbilical view, Sample KD20.

Figure 19- *Praemurica uncinata* (Bolli), Spiral view, Sample KD13.

Figure 20- *Morozovella angulata* (White), Side view, Sample KD19.

Figure 21- *Globanomalina compressa* (Plummer), Side view, Sample KD12.

Figure 22- *Globoconusa conusa* Khalilov, Spiral view, Sample KD6.

Figure 23- *Morozovella velascoensis* (Cushman), Umbilical view, Sample KD19.

Figure 24- *Morozovella conicotruncana* (Subbotina), Spiral view, Sample KD19.

PLATE II

