Foraminifers of the Late Cretaceous- Early Eocene in central Tunisia: Biostratigraphy and paleoenvironment

Oum elkhir Mahmoudi^{1.*}, Nebiha Ben Haj Ali¹

1- Département de Géologie, faculté des sciences de Tunis. 1060 El Manar II, Tunis, Tunisie.

* Corresponding Author: mahmoudioumelkir@hotmail.fr

Received: 28 February 2016 / Accepted: 25 December 2016 / Published online: 26 December 2016

Abstract

Biostratigraphic and statistical studies were carried out on the Late Cretaceous- Early Eocene succession of Khanguet Zalga in Jebel Serj area, central Tunisia. Paleontological analysis shows the occurrence of about 300 species (200 benthic foraminifera and 48 planktonic foraminifera). The most significant foraminifera (relative abundance more than 5%) are indicated in the vertical charts of distribution. Based on the vertical stratigraphic distribution of the planktonic foraminiferal species, six zones and subzones are recognized: Gansserina gansseri zone of Late Cretaceous (lower Maastrichtian), Igorina albeari (P3b) zone of Middle Paleocene (Selandian), Acarinina soldadoensis (P4c) subzone of the upper Paleocene, Morozovella velascoensis (P5) zones of Late Palaeocene (Thanetian), Morozovella edgari (E3) and Morozovella subbotinae (E5) zones of Early Eocene (Lower Ypresian). Biotic parameters and benthic assemblages from Late Cretaceous- Early Eccene Serj section allow the reconstruction of paleoenvironment and paleoecological fluctuations. In fact, our records indicate that the environment evolved from an initially oligotrophic, middle bathyal depositional environment for Gansserina gansseri, and P4c zone to upper bathyal setting for P3b and P5 towards a more eutrophic inner neritic setting for E3 and E5. The high abundance and the low diversity in this section are related to the sedimentological, the minimum of dissolved oxygen and the paleodepth conditions. The ecosystem is unbalanced for middle bathyal environments (hyaline shell-groups), balanced for shallow environment and upper bathyal (agglutinated and calcareous shell-groups).

Keywords: Central Tunisia; Biostratigraphy; Paleo-diversity; Paleoenvironment; Foraminifera.

1- Introduction

The Paleocene -Early Eocene interval has been studied by several authors. It includes two spectacular events: the Cretaceous-Tertiary (K/T) boundary (characterized by a massive planktonic foraminifera extinction; Salaj, 1980; Keller, 1988; Coccioni and Galeotti, 1994; Keller *et al.*, 1998 and Gearty, 2013) and Paleocene/Eocene boundary (Benthic Extinction Event (BEE); Tjalsma and Lohman, 1983; Thomas, 1990; Berggren and Aubry, 1996; Zili *et al.*, 2009). The global warming event, recorded during the Paleocene and culminating in the Early Eocene, represents an important phase of biotic and environmental change (Zachos *et al.*, 2001, 2003; Takeda *et al.*, 2007).

In Tunisia, Paleocene marl-shale hemipelagic series, with a diversified paleobiodiversity deposited in open marine conditions, have been influenced by terrigenous input from Kasserine Island and Algerian promontory (Bishop, 1985; Zaier et al., 1998). Undoubted gaps in sedimentation, recorded in the Paleocene of eastern Tunisia, indicate the action of synsedimentary movements leading to some short-term shallowings or even local emergence which is confined to the areas where sedimentation was related to a so-called "hautfond" phenomenon. The Eocene corresponds to a prolific period for the development of Nummulite and planktonic foraminifera carbonate platforms along the continental margins of the Tethyan Ocean (Zaier *et al.*, 1998; Jorry, 2003).

The main purpose of our research was to accurately correlate Late Cretaceous- Early Eocene of the central Tunisia with the Standard biostratigraphic scales. Moreover we discuss for the first time the relationship between paleodepth, morphology and shells groups of benthic foraminifera. Finally we observe if the global warming events have impact on the quantitative distribution of foraminifera.

In this paper we compile multiple sources of bio-magneto- astro-chronological time scale like Loeblich and Tappan, 1957; Berggren, 1960; Blow and Banner, 1962; Blow, 1979; Berggren 1995; Berggren and Pearson, 2005; Wade *et al.*, 2011.

2- Geological Setting

The study area "Khanguet Zalga" is part of the southern edge of the Serj–Bargou massif from the Tunisian Central Atlas (Fig. 1). This anticlinal localizes 3 km from the city of Oueslatia. It is far 25 km South-East from the capital of Seliana. The main geological formations encountered in the section are the EI Haria bounded by the upper beds of the Abiod Formation (upper Maastrichtian) below and those of the Bou Dabbous (Eocene) above. Undoubted gaps in sedimentation, recorded in the Paleocene of Eastern Tunisia, indicate the action of synsedimentary movements. The section was chosen in view of its expanded and well exposed nature.



Figure 1) Structural framework and facies distribution of the Eocene in the region of Oueslatia (According Jorry et al. 2001, as amended).

3- Material and methods

The studied material is extracted from 116 samples collected from the Khanguet Zalga section. Samples are spaced from 10 cm to 1 m within for shaley and marly sediments and 30 cm to 10 m in massive calcareous beds. About 100 grammes of each sample were treated with a 10% hydrogen peroxide solution during a day. After disintegration, samples were washed through a sieve column metallic background with mesh sizes decreasing from top to bottom (2 mm to 0.63 μ m) in order to remove the organic and clay fractions. The dry residues were sieved into three fractions (>63 μ m, >250 μ m, and >500 μ m). Each fraction was examined binocular microscope. All under the foraminifera observed were picked, identified, counted, MEB-photographed and placed in foraminifer's slides and covered for safety and future reference. Most of species are illustrated on Plates 1 through 4. The slides were properly labelled with well name and sample reference. The foraminiferal species was determined on the base of several publishing like Ellis and Messina (1940) and Loeblich Tappan's (1988).

In the discussion below we differentiate between lowest (LO) and highest (HO) occurrences of marker species used to define biozone boundaries. Reference sources for the zonal datum's that have been paleomagnetically calibrated and their ages are presented in Figure 2. We adopt the naming convention for five categories of interval zones as described by Berggren and Pearson 2005 and Wade *et al.* 2011.

In order to define paleoenvironmental conditions, 24 samples were selected for this study. The multivariable statistical analyses based on a wide range of diversity indexes (specific richness RS, Shannon-Weaver index HS (Murray 1991), dominance index D (den Dulk 2000), Fisher α index (Williams 1964) and equitability E (Murray 1991) are so useful for paleoenvironmental interpretations.

4- Results

4.1- Lithology and micropaleontology analysis

From the base to the top we have four lithological units in the Late Cretaceous- Early Eocene interval of Khanguet Zalga section.

The upper beds of the Abiod Formation (4 m thick): This formation is the oldest exposed rock unit in the study area. It consists fossiliferous of whitish grey limestone packstones with same quartz and dolomite crystal and few beds marly limestones intercalations; the washed marls contains many forms such as the Globotruncanidae with, Kuglerina rotundata, Rugoglobigerina reicheli, Planomalina prairichillensis, Laeviheterohelix glabrans, Laeviheterohelix sp., Planomalina alvarez and Hedbergella monmouthensis for planktonic species. Benthic foraminifera are present throughout the set and delivered the forms: Lenticulina following rotulata, Dentalina reussi and Bolivina incrassata giganta. This set is attributed to lower Maastrichtian, Ganserina ganseri Zone. The top of this unit is topped by a hard ground area.

El Haria Formation (73 m thick): the base of this unit comprises dark-grey shales and marls with a gypseous vein intercalations and celestibarite nodule. Succeeded by green-grey shales and marls with nine thin beds of limestones. This unit shows many forms (Fig. 2 and 3) such as: Subbotina, Morozovella, Globanomalina and Acarinina for planktonic Nodosaria, genus and *Lenticulina*, Anomalinoides, Cibicidoides. Spiroplectammina, Bathysiphon, Hyperammina, Bulimina and Bolivina for benthic genus. The use of planktonic foraminiferal zonation in regional biostratigraphic studies has allowed for

dividing this set on three Zones. After the remaniement, appears the first index species *Morozovella angulata* concomitant with *Igorina albeari*, 2 m above the index species *Globanomalina pseudomenardii* have been

occurred. This part of the Formation corresponds to the subzone P3b of the Zone Morozovella angulata P3. It is interpreted as an of the Selandian. equivalent In this biostratigraphic interval P3b, the enrichment by agglutinated benthic foraminifers assemblage with same calcareous forms (Fig. 2 and 3), indicate that the climate became cooler compared to the same during the Late Cretaceous period (Sliter and Backer, 1972; Thomas, 1990a; Galeotti and Coccioni, 2002; Guasti et al., 2007). The Midway assemblage of Nodosaria-Dentalina (Fig. 3) indicates a water depth of about 400 to 500 m.

From the first occurrence of Acarinina soldadoensis (Fig. 2), simultaneously with two planktonic species datum Globanomalina pseudomenardii and Morozovella velascoensis, to the highest occurrences of Globanomalina pseudomenardii, 34 m thick is attributed to the subzone P4c of the Zone P4. Planktonic foraminifers generally predominate (Fig. 4), being represented by genera and species typical of open warm-water marine basin. Benthic foraminiferal assemblages are dominated by Bolivina, Nuttallides truempyi, Osangularia, Oridorsalis. Lenticulina. Anomalinoides and Cibicidoides (Fig. 3). The species present in this assemblage are characterized by moderated geographic distribution as they are known almost exclusively from deep sea sediments (Tjalsma and Lohmann, 1983; Hulsbos, 1986; Nomura, 1991; Thomas, 2003) penetrated by DSDP in Indian and Pacific oceans and known on land from the Transcaspian area, Italy and southern France. Summarizing analysis obtained on foraminiferal distribution in the subzone P4c Acarinina soldadoensis/Globanomalina pseudomenardii show a middle bathyal depositional environment, with a paleodepth included between 500-600 m.

The top of this Formation is attributed to the later zone of Paleocene *Morozovella velascoensis* (P5) (Fig. 2). It is characterized by the partial range of the nominate taxon between

the highst occurrences of Globanomalina pseudomenardii and the first occurrence of Morozovella edgari. In this zone. two depositional environments were identified: the first is similar to the P4c, so middle bathyal depositional environment. The second is characterized by a decreasing contribution of the elements of the 'Velasco type' with mass occurrence of Midway Benthic foraminiferal assemblages (Fig.3), dominated bv the Anomalinoides welleri - cibicidoides dayi assemblage indicating a water depth of about 400m.

Chouabine Formation (1m thick): intervals marked by phosphatized marl clay containing shark and fish teeth and coprolites, but macro-vertebrate remains have not been observed. The Eocene period still with this unit, we have the first occurrences of index species Morozovella edgari and Morozovella marginodendata which are attributed to the Partial-range zone of the Morozovella marginodentata (E3) of the Early Ypresian. The most benthic species are Anomalinoides acuta, crosswicksona, Lenticulina Planularia coegaensis, Anomalinoides midwayensis and Lenticulina sp. and many large Frondicularia phosphatica (Fig. 3) associated to numerous fragments of brachiopod and gastropod shells, indicating periods of shallowing marine basin.

- Bou Dabbous Formation (23m thick): The base consists of black massive limestones with some marly intercalations which consist of diverse fauna planktonic forms indicate a foraminiferal Zones *Morozovella subbotinae* (E5) (Fig.2), this carbonate member is topped by a speckled grey hard massive limestone with *Globigerine* and few *Nummulite*. From the *Morozovella marginodentata* (E3) to the *Morozovella subbotinae* (E5) Benthic and planktonic foraminifers are relatively moderate (Fig. 4).



Figure 2) Planktonic species distribution of the Serj section.

UPPER- CRETACEOUS		PALEOCENE		EOCENE	ЕРОСН
MAASTRICHTIAN	SELANDIAN	THANETIAN		YPRESIAN	
			Rock		
Reusella insigui Gaudryina rugo Dentalina reuss	BENTHIC FORAMINIFERS				
Nodosaria- Dentalina		Nuttallides truempyi- Hperammina gaultia-Bathysiphon sp.	Cibicidoide s dayi - Anomalinoi des welleri	Anomalinoides acuta Cristellaria midwayensis	ASSEMBLAGES

Figure 3) Benthic species distribution of the Serj section.

4.2- Relationship between paleo-diversity and paleo-depth

4.2.1- Total microfauna

During the Paleocene-Early Eocene there was a general increase in foraminiferal densitie for all size fractions (Fig. 4). For the Early Cretaceous (Abiod Formation) we have the lowest numbers of microfauna compared to Paleocene - Early Eocene, there was a general decrease foraminifers presented a minimum density of ~120 specimens in Zel 15. The Paleocene (El Haria Formation) is characterized by the highest densities of microfauna individuals; they were recorded in Zel 53 and Zel 55 (more than 450 fossiliferous individuals). Figure 5 show a progressive dominance of benthic forms from the Early to Late Thanetian. During the Ypresian (Chouabine and Bou Dabbous formations) we have a clear decrease of the total microfauna with the equitability between benthic and planktonic forms. Based on the relative numbers of different shells types of benthic foraminifera we can divide the section in to three shells-zones.



Figure 4) Percentage of foraminiferal faunas.

From Zel9 to Zel35 (top of the Abiod and the base of El Haria Formation) we have the dominance of hyaline with same calcite shells (planktonic forms with Nodosaria and Lenticulina), between Zel 36 and Zel 72 (middle of El Haria Formation) we have the predominance of agglutinated shells (*Hyprerammina* and *bathysiphon*) the end (top of El Haria, Chouabine, and Bou Dabbous Formations) of the section is particularly rich in calcareous species (*Anomalinoides*, *cibicidoides* and lenticulina).

The ratio between planktonic and benthic foraminifera P/B has been calculated in order to

estimate paleodepth and productivity fluctuations. The results in figure 5 indicate that the environment evolved from an initially oligotrophic, deep sea, middle bathyal, to upper bathyal setting towards eutrophic outer neritic setting.



4.2.2- Generic diversity of benthic foraminifera

foraminifera Benthic are an important component of meiofaunal communities in many benthic environments, making them particularly useful for ecological as well as paleoecological studies. Table 1 presents the number of all genus that are >5% of all size fractions (500, 250 and 63μ m); it suggests the presence of three main faunal boundaries. Between Zel 9 and Zel 51 we have a phase of increase in number genus. Between Zel 53 and Zel 92 we have the maximum of diversity. Between Zel96 and Zel 108 there is a clearly decreasing of diversity.

Genus richness varies between the samples, the highest genus numbers were found at the thanetian samples (Zel 51 to Zel 96). Genus richness is also comparatively low in Zel 9. Generally, the genus number shows the same trends as the total density of biomass. The profile of Shannon index (Fig. 6) increases from the base to the top of the section. Values are less than 3 which indicate a minimum of dissolved oxygen concentration. The low values of fisher index reflect a low diversity and high dominance of same genus.

The evenness J (Fig. 6) has two different phases. The first consists of simples with values less than 0,8 (Zel 20, 25, 33, 40, 43, 46 and 48). It characterizes unbalanced assemblages with the biotope. When the evenness is upper to 0,8 the assemblages are stable and the environment is balanced. The dominance index D values (Fig. 6) are more than 0,1 indicate low diversity.

Throughout the section, the dissolved oxygen and the diversity are minimal from the bottom to the top, whereas the dominance is higher. The ecosystem is unbalanced for the deep and middle bathyal environments (hyaline shellgroups), balanced for shallow environment and

upperbathyal (agglutinated and calcareous shellgroups).

Table 1)) Abundance o	f Genus o	f benthic	foraminifera	(more than 5% d	of the total).
----------	---------------	-----------	-----------	--------------	-----------------	----------------

							U		0		0		0				0		,			
	Hypercountes	Bathyniphon	Annodiscus	Dentalina	Nodonaria	Netflabelinz	Lenticuâne	Saracenaria	Marginuispiris	Patradine	Marginailna	Boštrina	Bathrina	Anomalinolde	Gavelineita	clbicidnides	Gandyina	Spiroplectanina	Nettallder	Planstaria	Cristellaria	Virgalina
Zel 9	0	0	0	32,20	28,81	38,98	0	0	0		0	23		0	0	0	0	0	0	0	٥	0
Zel 15	0	0	0	8,19	13,11	22,95	6,55	8,19	11,47	18,03	11,47	0	5	7	0	0	11	7	0	0	0	0
Zel 20	0	0	0	5,08	15,25	10,16	52,54	1,69	15,25	0				9	0	0	0	0	0	0	0	0
Zel 25	0	0	0	5,67	30,49	41,13	2,12	7,80	5,67	7,09	0	11	8	10	0	0	0	0	0	0	0	0
Zel 29	0	0	0	5,88	30,39	51,96	3,92	4,90	2,94	0	0	4	5	3	0	0	0	0	0	0	0	0
Zel 33	0	0	0	6,03	25	49,13	0,86	5,17	6,89	6,89	0	6	8	8	0	0	0	0	0	0	0	0
Zel 36	15,33	8,66	2,66	8	9,33	11,33	14	8,66	12,66	9,33	0	17	21	0	0	0	13	19	14	0	0	0
Zel 38	13,23	8,08	0,73	15,44	5,14	9,55	11,76	18,38	12,5	5,14	0	16	25	0	0	0	0	17	7	0	0	0
Zel 40	26,48	28,97	0,62	5,29	11,83	2,18	6,54	1,86	1,55	1,55	1,85	7,78	2,80	0,62	0	0	0	9	2	0	0	0
Zel 43	34,43	31,78	2,31	1,32	3,64	1,98	4,63	0,99	1,98	0,66	2,64	4,30	5,62	3,64	0	17	0	11	0	0	0	0
Zel 46	7,352	2,45	0,49	1,47	27,94	2,45	31,37	3,43	3,43	0,49	4,41	8,33	6,37	0		13	0	0	0	0	0	0
Zel 48	25,70	29,09	1,97	1,69	3,38	1,97	5,93	0,56	1,41	0,84	3,10	7,06	6,21	7,90	1,41	1,69	0	6	0	0	0	0
Zel 51	20,95	6,287	2,69	4,79	12,57	1,49	16,16	2,39	5,08	1,19	4,49	6,28	4,79	6,58	4,19	0	14	0	0	0	0	0
Zel 53	20,09	22,14	1,36	2,73	11,64	1,82	14,38	2,73	1,82	0,22	2,28	5,02	5,93	2,51	3,19	2,05	0	0	0	0	0	0
Zel 55	20,98	23,83	2,07	2,33	8,54	2,59	12,69	1,81	2,33	0,77	2,33	6,21	5,18	2,33	4,14	1,81	0	0	0	0	0	0
Zel 59	17,66	13,33	2,33	7,65	5,66	13,66	1,66	3,33	433	8,66	5,66	5	6,33	4,65	0	19	14	0	0	0	0	0
Zel 63	10,65	13,01	4,73	12,42	5,32	13,90	3,84	2,07	3,25	9,46	2,07	5,32	4,73	5,62	3,55	0	12	0	0	0	0	0
Zel 76	3,53	10,28	7,39	14,45	7,07	17,04	5,46	2,57	4,50	8,36	2,89	3,53	4,18	5,46	3,21	٥	10	0	0	0	0	0
Zel 85	0	0	0	7,19	9,93	6,50	10,61	5,13	2,39	6,16	10,61	4,79	12,67	6,50	11,64	5,82	0	0	0	0	0	0
Zel 92	0	0	0	6,57	12,45	7,95	14,53	6,22	4,15	5,53	11,56	4,15	9,68	4,15	9,68	3,11	0	0	0	0	0	0
Zel 96	0	0	0	5,98	8,09	5,63	15,49	5,28	3,52	6,33	5,63	4,92	16,19	2,11	7,39	2,46	10,91	0	0	31	0	0
Zel 161	0	0	0	0	23,71	37,11	16,49	22,68	0		0			0	0	0	0	0	0	22	0	0
Zel 168	0	0	0	0	14,90	25	12,98	17,78	13,46	15,86	0			0	0	0	0	0	0	37	28	33



Figure 6) Biodiversity index fluctuations.

5- Discussion

In the Serj section, the lack area which extends from *Gansserina gansseri* to *M. angulata* zone (*Igorina albeari* P3b subzone) covers 10 MA; this section appears to be the most completely log in the region. Since other surrounding sections (plateau of Kesra, Oued Bahloul and Touijine) have sedimentary gaps between 12 to 14MA (Lajnef *et al.*, 2005; Jorry, 2004).

The hiatus show a break in the sedimentation accompanied by previous erosion filed marls El Haria Formation. It may be explained by the presence gypseous vein intercalations and celestibarite nodule which indicate an uprising in the region during periods of Maastrichtian – Paleocene.

The benthic faunal assemblages in the studied section may reflect different depositional environments. From deep sea for the *Ganserina ganseri* zone and the *Igorina albeari* subzone P3b with a water depth of about 400–500 m, a middle bathyal depositional for the subzone P4c *Acarinina soldadoensis/ Globanomalina pseudomenardii* with a paleobathymetry included between 500-600 m, to a phase of shallowing from upper bathyal to outer neritic

setting for the end of the section. The paleoecological indexes of benthic foraminifera illustrate three gradual shell groups (hyaline, agglutinated and calcareous groups) with decreasing paleodepth and age. The high abundance and the low diversity in this section related to the sedimentological, are the minimum of dissolved oxygen and the paleodepth conditions. A positive relationship was observed between shell-groups, paleodepth and lithology. The paleodepth fluctuations in this region are related to the subsidence phenomena.

The minimum of dissolved oxygen have a negative impact, by limiting the diversity of individuals. Currents with a minimum of oxygen can explain this situation. Therefore, only species that tolerate oxygen deficiency persist. This is deduced from the low diversity and the high abundances of some epifaunal species.

We don't have clear relationship between the distributions of foraminifers and the Global events (K/T) and the Paleocene–Eocene thermal maximum (PETM). During the late Paleocene, both shallowing of the marine basin and sedimentary gap in Central Tunisia is common with the sections of The Boreal (Gradstein et al., 1988, 1992, 1994; Nagy, 2005) and the Meridional provinces in Europe (Szczechura and Pozaryska 1971; Gradstein and Bäckström, 1996). Several regions of province were recorded by sedimentary gaps and occupied by hyposaline or even fresh-water, usually occupying shallow basins with restricted marine microfauna (Akhmetiev et al., 2010; Oreshkina, 2012). Planktonic microfauna so important for accurate dating of deposits are rare or absent. All the European stratotypes of the Paleocene-Early Eocene were epicontinental, bounded by deposits representing transgressional or regressional phases. These sections are bounded by hard ground surfaces. The group of benthic forms primarily comprises elements of the

Velasco (Nuttallides truempyi, type **Osangularia** velascoensis, **Bulimina** trinitatensis...) typical of deep sea sediments of the Tethys. Their occurrence indicates the proximity of a deep marine basin. Therefore it may be concluded that this assemblage lived on deeper shelf close to the continental slope. A similar situation was found in the case of the Paleocene microfauna of Babica clays from the Polish Carpathians (Szczechura and Pozaryska, 1974) and elsewhere in Europe. There are, however, elements of the Midway type (Cibicidoides, Anomalinoides and Lenticulina), typical of an epicontinental shelf marine basin, of a rather pandemic type. They are known from the North America and southern and central Europe, and Australia (Liu and Olsson, 1992; Berggren and Norris, 1997; Hollis et al., 2003; Alegret and Thomas, 2005; Bown, 2005; Alegret, 2007; Alegretand Thomas, 2009; Hulland Norris, 2011 and Alegretet al., 2012). The Paleocene yields also some benthic elements unknown from the Boreal provinces, which are characteristic of the Tethyan province and the North African province (Frondicu nlaria phosphatica and Palmula sp.) caracterise the shallow deposit of the end Paleocene.

6- Conclusions

Micropaleontological study planktonic of foraminifera retrieved from Khanguet Zalga sections spanning the Late Cretaceous- Early interval Eocene vielded six planktonic foraminiferal zones: Gansserina gansseri zone of Late Cretaceous (lower Maastrichtian), Igorina albeari (P3b) zone of Middle Paleocene (Selandian), Acarinina soldadoensis (P4c) subzone of the upper Paleocene, Morozovella velascoensis (P5) zones of Late Palaeocene (Thanetian), Morozovella edgari (E3) and Morozovella subbotinae (E5) zones of Early Eocene (lower Ypresian).

Biostratigraphic and quantitative distribution of benthic and planktonic foraminifera allow the

reconstruction of sea-level and paleoenvironmental conditions. Our records indicate that the environment evolved from an initially deep sea sediments relatively poor benthic forms: especially in planktonic foraminifera are rich and diverse. After the sedimentary gap, Selandian marl clay is composed especially of midway warm-water forms, indicate an upper bathyal deposition. During the Thanetian we have two changements in paleodepth, from the middle bathyal for the base of planktonic foraminifera Zone P4c to upper bathyal for the Late of P4c to P5, these fluctuations are reflected by the extinctions of numerous species Velasco type like Nuttallides truempyi and Hyperammina gaultia. The lack of any clearly cold-water elements and the occurrence of other warm-water forms such as Buliminids suggest some increase in the temperature of the water. The Early Eocene period still with phosphatized marl clay with faunas dominated by shallow-marine taxa (Anomalinoides. Buliminids. Haplophragmoides). The top of the section is interpreted as indicating an outer shelf setting.

These micropaleontological analyses show that the section maybe one of the most completely section in the region.

Acknowledgements

The authors thank Professor Rodolfo Coccioni, for useful advices and help, as well as reviewers for their collaboration and revisions, which greatly improved this manuscript.

References

- Akhmetiev, M. A., Zaporozhetz, N. I., Iakovleva, A. I. 2010. Comparative analysis of marine Paleogene sections and biota from West Siberia and the Arctic ocean. Stratigraphy and Geological Correlation: 18, 635-659.
- Oreshkina, T. V. 2012. Evidence of Late Paleocene- Early Eocene hyperthermal

events in biosiliceous sediments of Western Siberia and adjacent areas. Austoralian Journal of Earth Sciences: 105, 145–153.

- Alegret, L., Thomas, E. 2005. Cretaceous/Paleogene boundary bathyal paleo-environments in the central North Pacific (DSDP Site 465), the Northwestern Atlantic (ODP Site 1049), the Gulf of Mexico and the Tethys: the benthic foraminiferal record. Palaeogeogr. Palaeoclimatol. Palaeoecol: 224, 53-82.
- Alegret, L., Thomas, E. 2007. Deep-sea environments across the Cretaceous/ Paleogene boundary in the eastern South Atlantic Ocean (ODP Leg 208, Walvis Ridge). Marine Micropaleontology: 64, 1– 17.
- Alegret, L., Thomas, E. 2009. Food supply to the sea floor in the Pacific Ocean after the Cretaceous/Paleogene boundary event. Marine Micropaleontology: 73, 105–116.
- Alegret, L., Thomas, E., Lohmann, K. C. 2012. End-Cretaceous marine mass extinction not caused by productivity collapse. Proceeding of National Academy of Sciences: 109, 728– 732.
- Alegret, L., Thomas, E. 2013. Benthic foraminifera across the Cretaceous/ Paleogene boundary in the Southern Ocean (ODP Site 690): Diversity, food and carbonate saturation. Marine Micropaleontology: 105, 40–51.
- Berggren, W. A., Kent, D. V., Swisher, C. C., Aubry, M. P. 1995. A revised Cenozoic geochronology and chronostratigraphy, in Berggren, W.A., Kent, D.V., Hardenbol, J. (eds.), Geochronology, Time Scales and Global Stratigraphic Correlations: A Unified Temporal Framework for an Historical Geology. Society of Economic Paleontologists and Mineralogists, Special Volume: 54, 129–212.

- Berggren, W. A., Pearson P. N. 2005. A revised tropical and subtropical Paleogene planktonic foraminiferal zonation. Journal of Foraminiferal Research: 35, 279–298.
- Blow, W. H. 1979. The Cainozoic Foraminiferida, v.I and 11. E. J. Brill, Leiden.1413 pp.
- Coccioni, R., Galeotti, S. 1994. K/T boundary extinction: Geologically instantaneous or gradual event? Evidence from deep-sea benthic foraminifera. Geology: 22, 770–782.
- Den Dulk M. 2000. Benthic Foraminiferal Response to Late Quaternary variations in Surface Water Productivity and Oxygenation in the Northern Arabian Sea. Universiteit Utrecht, Utrecht, 205 pp.
- Galeotti, S., Coccioni, R. 2002. Changes in direction of Cibicidoides coiling (Nakkady) pseudoacutus across the Cretaceous-Tertiary boundary of Tunisia: palaeoecological and biostratigraphic implications. Palaeogeography, Palaeoclimatology, Palaeoecology: 178, 197-210.
- Gradstein, F. M., Agterberg, F. P., Aubry, M.-P., Berggren, W. A., Flynn, J. J., Hewitt, R., Kent, D. V., Klitgord, K. D., Miller, K. G., Obradovich, J. Ogg, J. G., Prothero, D. R., Westermann, G. E. G. 1988. Chronology of fluctuating sea levels since the Triassic a critique. Science: 235, II56–II67
- Gradstein, F. M., Agterberg, F. P., D'Iorio, M.A. 1988. Time in Quantitative Stratigraphy.Proceedings Workshop on Quantitative Dynamic Stratigraphy, Denver, Colorado, Prentice Hall, p.519–543.
- Gradstein, F. M., Kaminski, M. A., Berggren,
 W. A. 1988. Cenozoic foraminiferal biostratigraphy of the Central North Sea; in Rögl, F. and Gradstein, F.M. (eds.). Proc. Conference Agglutinated Benthic Foraminifera, Vienna 1986, p. 97–108.

- Gradstein, F. M., Von Rad, U. 1992. Stratigraphic evolution of Mesozoic continental margin and oceanic sequences, NW Australia and N. Himalayas. Marine Geology: 74, 123–152.
- Gradstein, F. M., Kaminski, M. A., Berggren,W. A., Kristiansen, I. L., D'Iorio, M. 1994.Cenozoic biostratigraphy of the North Sea and Labrador Shelf. Micropaleontology,Special Publication: 40, 152 p, 22 plates.
- Gradstein, F. M., Backstrom, S. 1996. Cenozoic biostratigraphy and paleobathymetry of Haltenbanken and northern North Sea. Norsk Geologisk Tidsskrift: 76, 3–32.
- Guasti, E., Speijer, R. P. 2007. The Paleocene-Eocene thermal maximum in Egypt and overview of the planktic Jordan: an foraminiferal record. In: Monechi, S., Coccioni, R. and Rampino, M. (Eds), Large ecosystem perturbations: causes and consequences. GSA Special Publication: 424, 53-67.
- Hammer, Ø. D. A. T, Ryan, P. D. 2001. PAST
 Paleontological statistics software package
 for education and data análisis.
 Palaontologica Electronica: 4, 9 pp.
 (http://palaeoectronica.org/2001_1/past/issu1_01.htm).
- Hollis, C. J., Strong, C. P., Rodgers, K. A., Rogers, K. M. 2003. Paleoenvironmental changes across the Cretaceous/Tertiary boundary at Flaxbourne River and Woodside Creek, Eastern Marlborough New Zealand. New Zealand Journal of Geology and Geophysics: 46, 177–197.
- Hulsbos, R. E. 1986. Eocene benthic foraminifers from the upper continental rise off New Jersey, deep sea drilling project site 6051.
- Jorry, S. 2004. The Eocene nummulite carbonates (central Tunisia and NE Libya) sedimentology, depositional environments,

and application to oil reservoirs. These de 3ème cycles.

- Keller, G. 1988. Extinction, survivorship and evolution of planktic foraminifera across the Cretaceous/Tertiary boundary in EI Kef, Tunisia. Marine Micropaleontology: 13, 535–551.
- Lajnef, B., Le Callonnec, L., Yaich, C., Renard, M., Benzarti, R. 2005. L'intervalle Paléocène supérieur – Eocène inférieur sur un profil de la Tunisie centro-septentrionale : approches sédimentologique et chimiostratigraphique. Bulletin de l'Institut Scientifique, Rabat, section Sciences de la Terre.
- Liu, C., Olsson, R. K. 1992. Evolutionary adaptive radiation of microperforate planktonic foraminifera following the K/T mass extinction event. Journal of Foraminiferal Research: 22, 328–346.
- Loeblich, A. R. Jr., Tappan, H. 1957. Planktonic foraminifera of Paleocene and early Eocene age from the Gulf and Atlantic Coastal Plains. Studies in foraminifera: Part 1 Planktonic foraminifera. United States National Museum Bulletin: 251, 191 p.
- Murray J.W. 1991 a. Ecology and distribution of benthic foraminifera. In: Lee, J.J., Anderson, O.R. (Eds.). Biology of Foraminifera. Academic Press, pp. 221–254.
- Murray, J. W. 1991 b. Ecology and paleoecology of benthic foraminifera. Longman, 398 p.
- Nagy, J. 2005. Delta-influenced foraminiferal facies and sequence stratigraphy of Paleocene deposits in Spitsbergen. Paleogeography, Paleoclimatology, Paleoecology: 222, 161–179.
- Nomura, R. 1991. Paleoceanography of Upper Maastrichtian to Eocene benthic foraminiferal assemblages at sites 752, 753, and 754, Eastern Indian Ocean, in Peirce, J., Weissel, J., Taylor, E., Alt, J., Dehn, J.,

Driscoll, N., Farrell, J., Fourtanier, E., Frey, F., Gamson, P. D., Gee, J. S., Gibson, I. L., Janecek, T., Klootwijk, C., Lawrence, J. R., Littke, R., Newman, J. S., Nomura, R., Owen, R. M., Pospichal, J. J., Rea, D. K., Resiwati, P., Saunders, A. D., Smit, J., Smith, G. M., Tamaki, K., Weis, D., Wllkinson, G. (eds.), Broken Ridge and Ninetyeast Ridge: Proceedings of the Ocean Drilling Program, Scientific Results: 121, 3– 29.

- Pearson, P. N. 1995. Planktonic foraminifer biostratigraphy and the development of pelagic caps on guyots in the Marshall Islands Group. In: Haggerty J., Premoli Silva, I. Rack, F. McNutt M.K. (Eds.). Proceeding of the Ocean Drilling Program: Scientific Results: 144, 21–59.
- Plummer, H. J. 1927. Foraminifera of the Midway Formation in Texas, University of Texas Bulletin: 2644, 1–201.
- Reali, S., Ronchi, P. 1998. Sedimentological model of nummulitic limestone, El Garia Formation (Central Tunisia and offshore Libya). Tertiary to Recent Larger Foraminifera. Conference-Workshop- Fiel Trip, Kingston, UK, 41-42. Reiss, Z. and Hottinger, L., 1984.
- Salaj, J. 1980. Microbiostratigraphie du Crétacé et du Paléogène de la Tunisie Septentrionale et Orientale (Hypostratotypes Tunisiens). Bratislava, Slovakia, 238 p.
- Sliter W. V., Baker R. A. 1972. Bathymetric distribution of benthic foraminifers. Journal of Foraminiferal Research: 2, 167–183.
- Szczechura, J., Pozaryska, K. 1971. The Montian warm-water foraminifers in the Meridional Province of Europe. Acta Palaeontologica Polonica: 16, 345–388.
- Pozaryska, K., Brochwicz-lewirski, W. 1974.
 Wspolczesne ujecie zagadnienia prowincji paleogeograficznych. Postqpy Nauk Geol., 6, 546. Szczechura, J. and Pozaryska, K.

Foraminiferida from the Paleocene of Polish Carpathians (Babica Clays). Acta Palaeontologica Polonica: 31, 1–142.

- Takeda, K., Kaiho, K. 2007. Faunal turnovers in central Pacific benthic foraminifera during the Paleocene-Eocene thermal maximum: Palaeogeography, Palaeoclimatology, Palaeoecology: 251, 175–197.
- Tjalsma, R. C., Lohmann, G. P. 1983. Paleocene-Eocene bathyal and abyssal benthic foraminifera from the Atlantic Ocean. Micropaleontology, Special publication: 4, 90 p.
- Thomas, E. 1990, Late Cretaceous-early Eocene mass extinctions in the deep sea, in Sharpton, V. L., D. Ward, P. (eds.), Global Catastrophes in Earth History: An Interdisciplinary Conference on Impacts, Volcanism, and Mass Mortality: Geological Society of America, Special Paper: 247, 481-495.
- Thomas, E. 2003. Extinction and food at the sea floor: A high-resolution benthic foraminiferal record across the Initial Eocene Thermal Maximum, Southern Ocean Site 690. In: Wing, S. L., Gingerich, P. D., Schmitz, B., and Thomas, E., Eds. Causes and Consequences of Globally Warm Climates in the Early Paleogene. Boulder, Colorado. Geological Society of America Special Paper: 369, 319–332.

- Wade, B. S., Pearson, P. N., Berggren, W., Pälike, H. 2011. Review and revision of Cainozoic tropical planktonic foraminiferal biostratigraphy and calibration to the geomagnetic polarity and astronomical time scale. Earth-Science Reviews: 104, 111–142.
- Zachos, J. C., Pagani, M., Sloan, L., Thomas, E., Billups, K. 2001. Trends, rythyms, and aberrations in global climate 65 Ma to Present: Science: 292, 686–693.
- Zachos, J. C., Wara, M. W., Bohaty, S., Delaney, M. L., Petrizzo, M. R., Brill, A., Bralower, T., Premoli-Silva, I. 2003. A transient rise in tropical sea surface temperature during the Paleocene-Eocene Thermal Maximum: Science: 302, 1551– 1554.
- Zaier, A., Beji-Sassi, A., Sassi, S., Moody, R. T.
 J. 1998. Basin evolution and deposition during the early Paleogene in Tunisia.In: MacGregor, D. S., Moody, R. T. J., Clark-Lowes, D. D. (Eds.). Petroleum Geology of North Africa. Geological Society, Special Publication: 375–393.
- Zili, L., Zaghbib-Turki, D., Alegret, L., Arenillas, I., Molina, M. 2009. Foraminiferal turnover across the Paleocene/Eocene boundary at the Zumaya section, Spain: record of a bathyal gradual mass extinction. Revista Mexicana de Ciencias Geológicas: 26, 729–744.

Appendices:



Plate 1) 1: Neoflabellina munismalis (WEDEKIND) (Zel.2); 2: Dentalina reussi NEUGEBOREN (Zel.2); 3: Vaginulinopsis longiforma (PLUMMER) (Zel.2); 4: Nodosaria limbata d'ORBIGNY (Zel.10); 5: Frondicularia sp. (Zel.24); 6: Marginulina sp. (Zel.15); 7: Bulimina quadrata PLUMMER (Zel.16); 8: Marginulina sbbullata Hantk. (Zel.16); 9-10: Lenticulina rosetta (Zel.6); 11: Lenticulina degolyeri (PLUMMER) (Zel.16); 12-13: Lenticulina pilulifera (CUSHMAN) (Zel.15); 14: Lenticulina klagshmnensis (BORTZEN) (Zel.16); 15: Lenticulina sp. (Zel.16); 16: Ammodiscus cretaceous CUSHMAN et HAN (Zel.15); 17-18: Vulvulina pennatula (Zel.15); 19-20: Anomalina granosa (Zel.33).



Plate 2) 1: (Zel.38); 2-3: Lenticulina turbinata (Zel.35); 4: Sarasenaria tunesiana TEN DAM et SIGAL. (Zel.35); 5: Vaginulopsis midwayena (Zel.35); 6: Spicule d'échinoderme (Zel.38); 7: Nodosaria macneili CUSHMAN (Zel.38); 8: Marginulina tuberculata (PLUMMER) (Zel.38); 9: palmula toulmini (Zel.23); 10-11 : Cibicidoides praecursoria (SCHWAGER) (Zel.46); 12: Frondicularia phosphatica (Zel.63); 13-14: globulina gibba d'ORBIGNY (Zel.63); 15-16: Gyroidinoides subangulata (PLUMMER) (Zel.63).



Plate 3) 1-2-3: Pseudotextularia nuttalli (Zel.1); 4-5: Globotruncana bulloides (Zel.1); 6-7: Globotruncana falsostuarti (Zel.2); 8- 9-10-11-12: Globotruncana stuartiformis (Zel.4); 13-14-15: Planomalina alvarez Zel.4); 15-16: Globotruncana linneiana (Zel.4).



Plate 4) 1-2: Globanomalina pseudomenardii (Zel.48); 3-4: Eogobigerina edita (Zel.35); 5-6-7: Morozovella acuta (Zel.41); 8-9: Subbotina velascoensis CUSHMAN (Zel.47); 10-11: Subbotina sp.1 CUSHMAN (Zel.50); 12: Subbotina sp.2 (BECKMANN) (Zel.50); 13-14-15-16: Morozovella velascoensis (CUSHMAN) (Zel.56).