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On the late Pleistocene-Holocene sapropel record of climatic and oceanographic variability in the eastern Mediterranean

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Abstract. Sapropels and intercalated marls in a piston core from the eastern Mediterranean are chemically and mineralogically distinct: kaolinite, smectite, and total S contents, Fe/Al, Ba/Al, Co/Al, Mo/Al, and V/Al are higher, whereas quartz, Mg calcite, illite contents, Si/Al, Ti/Al, K/Al, Rb/Al, and Zr/Al are lower in the sapropels. Missing and "ghost" sapropels are identified by mineralogical and chemical properties that are not prone to diagenesis. Primary production was higher (Ba/Al) and bottom water and/or interstitial oxygenation was lower (Co/Al, Mo/Al, Ni/Al, and V/Al) during sapropel formation. Wind speeds (quartz/clay, Si/Al, and Zr/Al) and bottom water salinities (Mg calcite/calcite) were higher during periods of marl formation. Sapropels represent a fundamentally different sedimentary facies whose formation is linked to changes in the hydrological balance in the basin, driven by precessionally modulated changes in monsoon strength and subtropical precipitation changes, which altered circulation, production, and sediment source areas.

1. Introduction

Sequences of organic-rich muds (sapropels) in the Holocene and late Neogene coccolith-foraminiferal marl oozes of the eastern Mediterranean have been a rich source of information on palaeoclimatic and palaeoceanographic variability [Olausson, 1961; Williams and Thunell, 1979; Rossignol-Strick, 1985; Rohling, 1994]. These distinctive deposits were formed during periods of increased freshwater influx into the eastern basin when insolation-driven strengthening of the monsoon led to increased rainfall in the northern and southern Mediterranean borderlands. These influxes caused profound changes in surface and deep circulation, productivity, bottom water oxygenation, and sediment supply. The tight link between the intervals of sapropel deposition and insolation maxima at the precessional frequency (albeit with a lag of a few thousand years) led, beginning with the seminal paper of Hilgen [1987], to the extensive use of sapropel sequences in the circum-Mediterranean region for erecting high-resolution astronomical chronologies over the last 10 Myr [Hilgen et al., 1995; Langereis et al., 1997]. The recovery of sapropels in the western Mediterranean during Ocean Drilling Program Leg 161 [Murat, 1999] demonstrates that unusual changes in climatic and/or oceanographic conditions have intermittently affected the entire Mediterranean over the last 3 million years.

In a prescient paper, Bradley [1938] predicted the occurrence of laminated, organic-rich sediments in the glacial age sediments of the eastern basin of the Mediterranean on the basis of the likely change in climate and/or lowered sea level at this time. By analogy with hydrographically restricted fjords and the Black Sea he reasoned that a wetter and cooler glacial climate, as generally

accepted at that time [Brooks, 1949], would have reversed the circulation of the eastern basin because of the presence of a lower-salinity surface layer and this would have led to deep water stagnation due to restricted deep water renewal. Kullenberg [1952] and Olausson [1961] extended this reasoning after the sapropels were first sampled during the Swedish Deep-Sea Expedition, and they suggested that the increased supply of fresh water from the Black Sea and from the pluvial condition of the Mediterranean region after the glacial maximum were instrumental in creating an anoxic basin which led to the formation of the sapropels. This explanation would be consistent with the occurrence of many sapropels during "warming trends" [Ryan, 1972]. It is now known that the sapropels formed mainly after, not during, the glacial maxima and are closely related to the precessional cycle of the monsoons [Rossignol-Strick et al., 1982].

The evidence for the formation of the sapropels via increased organic carbon preservation under bottom water anoxia has recently been reviewed by Emeis and Party [1996] and Cramp and O'Sullivan [1999], and this mechanism is now widely accepted. Basin stagnation is held to be caused by greatly increased runoff [Rossignol-Strick et al., 1982; Rossignol-Strick, 1983, 1985] or an altered hydrographic balance which decreased evaporative water losses [Rohling and Hilgen, 1991; Rohling, 1994]. An additional factor originally suggested by Berger [1976] is a higher settling flux of organic matter caused by increased primary production when the circulation and the nutrient budget of the eastern basin were altered by the changed hydrographic conditions [Calvert, 1983]. Rohling [1989] suggested that pycnocline shoaling induced the formation of a deep chlorophyll maximum at times of sapropel formation, and Sancetta [1994] proposed that large fluxes of mat-forming diatoms were responsible for high carbon fluxes, a suggestion that has recently been confirmed for one of the prominent late Pleistocene sapropel

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horizons [Kemp *et al.*, 1999]. The current view appears to be converging on a combined mechanism, whereby high carbon fluxes cause oxygen drawdown in deep water, leading to enhanced organic preservation [Sancetta, 1999].

The palaeoclimatic and palaeoceanographic information stored in the sapropel records, whatever mechanism is principally involved in their formation, includes the sources of organic matter [ten Haven *et al.*, 1987a, 1987b, 1987c], changes in freshwater balance [Thunell and Williams, 1989; Rohling and Hilgen, 1991; Rohling and De Rijk, 1999], glacial-interglacial changes in atmospheric $p\text{CO}_2$ [Fontugne and Calvert, 1992], and relative nutrient utilization [Calvert *et al.*, 1992]. The goals of this study were to acquire further insight into environmental changes over several glacial-interglacial cycles in the eastern Mediterranean from a detailed consideration of the mineralogical and inorganic chemical compositions of the deposits, employing recent advances in the use of sedimentary tracers to reconstruct variable sediment sources, texture, redox changes, and palaeoproductivity.

2. Materials and Methods

Core MD 84641 (33°02'N, 32°38'E; 1375 m water depth) was collected during the *Noë* cruise of the RV *Marion Dufresne* in August, 1984. Separate samples were taken for foraminiferal isotopic and for chemical and mineralogical analyses. Sampling was at 10 cm intervals in the marl sections of the core and at 2 cm intervals through the sapropels. The foraminiferal isotope stratigraphy used here is taken from Fontugne and Calvert [1992]. Dried (60°C), powdered subsamples were used for mineralogical and chemical analysis. Organic carbon, total nitrogen and carbonate C determinations at the University of British Columbia (UBC) were carried out by a combination of

combustion/gas chromatography (total C and N) and coulometry (carbonate C); the precisions determined on replicate subsamples were $\pm 1.25\%$ and $\pm 3.5\%$ for total C and N and $\pm 2\%$ for carbonate. Organic carbon values were derived by difference with a combined precision of $\pm 2.35\%$. Organic C determinations at Gif-sur-Yvette were made by combustion followed by coulometric determination of the evolved CO_2 of subsamples that were soaked in 10% HCl overnight and dried to constant weight at 80°C. Absolute precision was better than 0.1%. Differences between the two organic carbon data sets were generally less than $\pm 5\%$. Separate subsamples were used for major and minor element determinations by X-ray fluorescence spectrometry, using methods described by Calvert [1990]. Precisions of the analyses were $\pm 3\%$ for the major elements and $\pm 5\%$ for the minor elements. The mineralogy of the bulk samples and the clay fraction ($< 2 \mu\text{m}$; separated by settling/centrifugation) was determined by conventional powder X-ray diffractometry. Relative abundances of clay, quartz, low-Mg calcite and high-Mg calcite in the bulk sediments were determined using the ratios of the peak intensities at 450 (common to all clay minerals), 426, 303, and 298 pm, respectively. Clay mineral abundances (smectite (ethylene glycol treatment), illite, kaolinite, chlorite, and palygorskite) of selected horizons within sapropels and above and below these intervals were determined from the ratios of their peak areas on X-ray diffraction (XRD) spectra. No significant variations in clay mineral crystallinity were detected.

3. Results and Discussion

3.1. Sapropel Stratigraphy

Core MD 84641 is located on the northern extremity of the Nile Cone between Cyprus and the Nile Delta. It consists of 11.61

Figure 1. Down core variation of organic carbon concentrations and the $\delta^{18}\text{O}$ (*Globigerinoides ruber*) stratigraphy of core MD 84641.

Figure 2. Age model for the core, showing a comparison of the $\delta^{18}\text{O}_{\text{ruber}}$ stratigraphy and the SPECMAP stack [Martinson *et al.*, 1987a] plotted against age, sapropel stratigraphy based on the age model for the core, and down core CaCO_3 variations. The SPECMAP curve, with events identified, has been offset to lighter values for ease of viewing the two isotope curves.

m of pale cream to yellowish brown nanofossil/foraminiferal marl ooze (hereinafter referred to as marl) with nine olive green sapropels varying in thickness from 6 to 22 cm (Figure 1). A hiatus at 445 cm depth is marked by a subtle change in the color and an abrupt change in water content. Four of the sapropels (those at 328–341, 374–393, 458–479.5, and 578–596 cm) contain gray mud layers with lower organic carbon contents, producing two maxima in organic carbon content at each level.

The chronostratigraphy of the core is based on the *Globigerinoides ruber* $\delta^{18}\text{O}$ record (Figure 1), ^{14}C dating of planktonic foraminifera in the upper 135 cm of the core, and palaeomagnetic reversal stratigraphy. The radiocarbon age of the uppermost sapropel is 6869–8570 years B.P. [Fontugne *et al.*, 1994] and is therefore S1 [Troelstra *et al.*, 1991]. The reservoir-corrected calendar ages [Bard *et al.*, 1998] are 7463 to 9491 years. The $\delta^{18}\text{O}_{\text{ruber}}$ record shows a range of -2.3 to $+3.2$ ‰, consistent with other foraminiferal records from the Mediterranean that all show a much larger glacial/interglacial range than that in the Atlantic [Vergnaud-Grazzini *et al.*, 1977]. Correlation with the SPECMAP stack [Imbrie *et al.*, 1984; Martinson *et al.*, 1987b] using the AnalySeries software [Paillard *et al.*, 1996] shows that the record in this core extends back to stage 12 (Figure 2). The $\delta^{18}\text{O}_{\text{ruber}}$ record suggests that the

hiatus at 445 cm represents 20 kyr of sedimentation [Kallel *et al.*, 2000]. The Last Glacial Maximum (stage 2) has $\delta^{18}\text{O}$ values higher than 3‰, whereas stage 8 has values close to 2‰. The lightest $\delta^{18}\text{O}$ values are found in stage 7.

Comparison of the organic carbon and $\delta^{18}\text{O}_{\text{ruber}}$ records (Figure 2) demonstrates that core MD 84641 contains sapropels S1 and S3–S10 [Cita *et al.*, 1977] or Si2, Si8–Si12, Si16–Si22, and Si30 using the proposed insolation-based numbering scheme of Langereis *et al.* [1997]. Sapropel S2 (Si4/6) appears to be missing, as is often the case [Rossignol-Strick, 1985] for this poorly developed interglacial stage. (Other missing sapropels in this core will be discussed subsequently.) The magnetic reversal Blake Event (beginning at 117 kyr B.P.) occurs between S4 (Si10) and S5 (Si12) [Tucholka *et al.*, 1987; Fontugne and Calvert, 1992], consistent with the isotope stratigraphy.

Apart from S6 (Si16), the sapropels occur in interglacial stages, there being three such horizons in stages 5 and 7, where there are three prominent interstadials. The grouping of sapropels into clusters has been noted previously by Hilgen [1990], who proposed that individual sapropels are directly related to minima in the precession index, and the clusters of sapropels are related to eccentricity. The organic-poor sections in sapropels S4 (Si10) and S8 (Si20) correspond with heavy $\delta^{18}\text{O}$ intervals in these warm

Figure 3. Relationship between organic carbon and total nitrogen concentrations in the core. The slope of the regression indicates the average ratio for the entire sample set; correction of the nitrogen values for the small amount of inorganic nitrogen present (shown by the intercept on the x axis) would produce a zero intercept, indicating that the organic C/N is roughly the same in the sapropels and the marls.

interstadials (Figure 2), emphasizing the close relationship between the occurrence of sapropels and climate in the eastern Mediterranean. In the case of S6 (Si16), $\delta^{18}\text{O}$ does not change across the organic-poor section.

The age model for this core yields sedimentation rates of between 1.3 and 5.3 cm/kyr. These are similar to those reported for this part of the eastern Mediterranean [Stanley and Maldonado, 1977]

3.2. Organic Carbon and Carbonate

Organic carbon contents of the sapropels range from 2 to 4.5 wt % (Figures 1 and 2). Intercalated gray mud layers with lower organic carbon contents in S4 (Si10), S6 (Si16), and S8 (Si20) produce two maxima in organic carbon content at each horizon. The marls have organic carbon contents ranging from 0.1 to 0.3 wt %, with no systematic trends in this lithology down the core. The $C_{\text{organic}}/N_{\text{total}}$ ratio is higher in the sapropels (16.92 ± 1.1), defined as those horizons with >1% organic C, compared with the marls (12.34 ± 4.1), a difference that is significant at the <1% level (two-sided t test). However, this difference is caused by the presence of a small amount of inorganic nitrogen in all samples, reflected in the intercept of the regression in Figure 3. Thus organic-poor samples have lower than average C_{organic}/N ratios, and organic-rich samples have higher than average C_{organic}/N ratios. Correction for the nitrogen fraction indicated by the intercept would produce a similar average C_{organic}/N ratio in both lithologies.

Calcium carbonate values have been estimated from the carbonate carbon determinations, assuming no phase other than calcite or aragonite is present (but see below). The concentrations range from 10 to 58% by weight and have minima just below or within all sapropels (Figure 2). The highest values occur in the upper part of stage 6, and this is followed by an abrupt change to the lowest values in the overlying sapropel S5 (Si12). High carbonate values also occur slightly above S1 (Si2), S7 (Si18), and S10 (Si30).

3.3. Mineralogy

Core MD 84641 contains variable proportions of quartz, feldspar, calcite, magnesian calcite, aragonite, smectite, illite, and kaolinite and minor chlorite, dolomite, and palygorskite. The

Figure 4. Down core distributions of the relative abundances of quartz and high-Mg calcite, expressed as the peak height ratios quartz/clay and high-Mg calcite/low-Mg calcite on X-ray diffractograms.

relative abundances of the principal phases show that the bulk composition of the sapropels and the marls differ significantly. Thus quartz/clay ratios are systematically lowest within and highest below each sapropel (Figure 4); furthermore, short-lived abrupt changes in this ratio occur throughout the core. The higher quartz contents of the marls signify that those intervals are coarser grained, whereas quartz/clay minima in the sapropels indicate that they consist of the finest grain sizes of the entire core.

In addition to high abundances of low-magnesian calcite, typical of foraminifera and coccoliths, there are also large changes in the abundance of high-magnesian calcite in the core (Figure 4). The spacing of the distinct d_{211} Mg calcite peak indicates that the Mg-rich phase contains ~16 mol% MgCO_3 [Goldsmith and Graf, 1958; Milliman and Müller, 1973]. Concentrations of this phase exceed those of low-Mg calcite in many of the marl sections of the core. The lowest values are found within the sapropels, where this phase is commonly undetectable using the powder diffraction method employed.

The sapropels have higher smectite and kaolinite and lower illite and palygorskite contents than the marls. Illite is the dominant clay mineral in the western part of the eastern Mediterranean, whereas smectite is dominant in the eastern part of the basin [Chamley, 1972; Dominik and Stoffers, 1978]. The high proportion of smectite in the clay fraction of core MD 84641 is attributed to its supply by the Nile [Venkatarathnam *et al.*, 1972; Stanley and Liyanage, 1986], and the higher content of kaolinite is probably due to its derivation from arid source areas in northern Africa [Foucault and Mélières, 2000]. Sapropels in the Bannock Basin in the central Ionian Sea also have much lower kaolinite/chlorite ratios (due to significantly higher chlorite

contents) and lower palygorskite contents than the associated sediments [Tomadin and Landuzzi, 1991]. Palygorskite characteristically forms in arid, saline soils that are typical of the circum-Mediterranean region [Singer, 1985]. It is presently transported over the central Mediterranean by winds blowing from southern sources [Tomadin *et al.*, 1984]. The lower abundance or the complete absence of this phase in the sapropels suggests that arid source areas were less extensive or wind strength was lower during times of sapropel formation.

3.4. Chemical Composition

Because the sapropels and marls in core MD 84641 have different and highly variable contents of organic matter and carbonate (Figure 2), which dilute the detrital fraction to variable extents, the compositions of the two facies have been compared by normalizing the major element contents to Al, a reliable index of fine-grained aluminosilicate detritus in terrigenous-bearing sediments [Calvert, 1976]. Minor element data from core MD 84641 support the conclusions that can be drawn from the major element variations, indicating that most of the Al is indeed contributed to the eastern Mediterranean within aluminosilicate detritus.

The down core distributions of Si/Al, Ti/Al, and K/Al are generally coherent, although Ti/Al behaves differently in some intervals (Figure 5). This coherence is consistent with the lack of significant biogenous opal in the core, a common observation in the eastern Mediterranean [Kemp *et al.*, 1999], which would otherwise cause variations in Si/Al ratios without affecting Ti/Al and K/Al. Minima in all three ratios characterize the sapropels, and there are also prominent minima in stages 8 and 9 and smaller relative minima in stages 3 and 6 where sapropels do not occur.

Figure 5. Down core distributions of Si/Al, Ti/Al, and K/Al. Ratios show distinct minima in all sapropels (apart from K/Al in S6) and between S1 and S3 and between S9 and S10.

Figure 6. Down core distributions of Mg/Al and Ca/Al ratios. The distributions are highly coherent, minima occurring in all sapropels except S6.

Sapropel S6 (Si16) has much higher Si/Al and K/Al ratios, and all three ratios are higher in the lower part of S8 (Si20) compared with the other sapropels. The major difference between the distributions of Si/Al and K/Al, on the one hand, and Ti/Al, on the other, is in the location of the highest values. For Si/Al and K/Al the maximal values are found below S5 (Si12) where Ti/Al ratios are very low, whereas the highest Ti/Al values are between S4 (Si10) and S5 (Si12).

These variations are probably due to bulk mineralogical changes, Si/Al varying with quartz content, K/Al varying with the abundance of K-feldspar and illite, and Ti/Al reflecting the relative abundance of Ti-bearing heavy minerals, such as ilmenite, sphene, etc. Si/Al and K/Al ratios are high where quartz/clay ratios are high (Figure 4). Thus, the Si/Al and K/Al ratios can be used as sensitive indices of the relative abundance of quartz and K-feldspar, respectively, and can be used as an indirect measure of sediment texture. The coarsest sediment (highest Si/Al and K/Al) occurs immediately below S1 (Si2), S5 (Si12), and S10 (Si30). Ti/Al ratios are low in stage 6 between S5 (Si12) and S6 (Si16), where Si/Al and K/Al ratios indicate the presence of coarse-grained material. The Ti-bearing heavy minerals may well be somewhat finer grained (although still coarser than the clay fraction) than quartz and feldspar, accounting for the differences in the locations of the Ti/Al and the Si/Al and K/Al peaks (Figure 5).

The distribution of Mg/Al ratios (Figure 6) is very similar to that of Si/Al and K/Al (Figure 5). Thus the values are lowest in all sapropels except S6 (Si16) and are highest in stages 2 and 6; prominent minima also occur in stages 8 and 9, where Si, Ti, and K are all low relative to Al. This could be due to the covariation of a Mg-bearing detrital aluminosilicate, such as chlorite or palygorskite, with quartz and feldspar. However, Mg-bearing

aluminosilicates, although present, are relatively minor constituents of the marls, and comparison of Figures 4 and 6 shows that the down core changes of high-Mg calcite and Mg/Al ratios are highly coherent. In view of the relative abundance of low- and high-Mg calcite (Figure 4) and the Mg content of the latter phase, it can be shown by calculation that the distribution of Mg is significantly controlled by the abundance of high-Mg calcite, which largely obscures any variability in aluminosilicate content. Down core trends in Ca/Al ratios are coherent with those of Mg/Al (Figure 6), reflecting the dominant control of the total Ca content by carbonate phases.

Fine-grained and nodular high-Mg calcite is common in the eastern Mediterranean [Olausson, 1961], where it occurs as roughly 6 mm diameter microcrystals in the marls but not in the sapropel horizons and also as nodules and cemented crusts on seamounts and on the deep basin floor [Fischer and Garrison, 1967; Milliman and Müller, 1973; Müller and Fabricius, 1974]. It is also abundant in cores from the Aegean Sea [Aksu et al., 1995]. Because of its abundance (up to 75 wt %) and mode of occurrence it is probably formed inorganically in the surface sediments [Milliman and Müller, 1973], and its precipitation in the eastern basin is due to the prevailing high bottom water salinities. This phase also occurs in the Red Sea [Gevirtz and Friedman, 1966; Milliman et al., 1969], where its presence has been attributed to high salinity conditions. Its occurrence in the marls in MD 84641 and its absence in the sapropels suggests that bottom water as well as surface water [Thunell and Williams, 1989] salinities were higher during the deposition of the marls. Other minima in the abundance of high-Mg calcite (Figure 4) where sapropels are absent and where Si/Al, K/Al, and Ti/Al are also low, notably in stages 3, 8, and 9, will be discussed in section 3.5.

Figure 7. Down core distributions of Fe/Al and wt % S. The latter occurs predominantly as pyrite.

The distribution of Fe/Al ratios (Figure 7) shows that Fe enrichments characterize many of the sapropels; however, S5 (Si12), the upper part of S6 (Si16), and S9 (Si22) have very small enrichments. Sapropel S10 (Si30) has higher Fe/Al in the lower part and very low values in the upper part. Much of the variation in the abundance of Fe in MD 84641 is probably controlled by the distribution of pyrite, as reflected in the distribution of total sulphur (Figure 7), which ranges from 0.01 to 4.35% by weight and is highest in the sapropels. There are also relatively high S concentrations just below many of the sapropels due to the downward diffusion of sulphide from the anoxic organic-rich horizons and its reaction with reduced Fe diffusing from deeper levels in the sediment [Passier *et al.*, 1999]. Relative Fe contents are much less variable in the marls, and Fe/Al is slightly lower in stages 2, 4, and 6, and the middle of stage 8, possibly indicating some variability in the nature of the detrital Fe-bearing aluminosilicate minerals in the core.

Variability in the minor element composition of the sapropels and associated sediments in core MD 84641 supports the interpretation of down core changes in bulk sediment composition and additionally provides information on changes in palaeoproduction and redox conditions. Thus the distribution of Sr closely follows that of Ca. Highest concentrations are found where CaCO_3 is highest, namely immediately below S5 (Si12) and S6 (Si16) and in stage 1, whereas the lowest Sr/Al values are found in the sapropels S5 (Si12) and S7 (Si18) (Figure 8). This reflects the distribution of biogenous calcite, in which Sr substitutes for Ca up to values of 1200–1500 ppm [Turekian, 1964; Thompson and Bowen, 1969], and of high-Mg calcite. Strontium concentrations in the latter, inorganic phase are likely to be lower than in biogenous carbonate phases, but we are not able to determine this partitioning because it was not possible to separate the very fine grained inorganic and biogenous (coccolith) materials. This conjecture is supported by the lack of significant

difference in the projected Sr content at 100% CaCO_3 in the sapropels and the marls.

Rubidium and Zr in the core are controlled entirely by the composition and grain size of the aluminosilicate fraction. Rubidium substitutes for K in most K-bearing minerals, so the distribution of these two alkali metals parallels each other down core (Figures 5 and 8); the lowest values are found in many of the sapropels, and the highest Rb/Al ratios occur in stages 2 and 5. Sapropels S6 (Si16) and S8 (Si20) show little change in both Rb/Al and K/Al. Zr/Al ratios change in concert with changes in Si/Al (Figures 5 and 8), and values of both ratios are positively correlated. Thus highest Zr/Al is found in stage 2 and the upper part of S6, and the lowest values occur in each of the sapropels. Zirconium occurs predominantly in the mineral zircon in marine sediments, which in turn, is normally confined to the fine-sand/coarse-silt fraction. Variations in Zr/Al therefore lend support to the inference from the quartz/clay and Si/Al ratios of larger grain sizes in the marl horizons below many of the sapropels.

Somewhat similar contrasts in Si/Al, Ti/Al and Zr/Al ratios between sapropels and marls have been identified in cores from Ocean Drilling Program (ODP) Sites 964 and 967 [Wehausen and Brumsack, 1999], and the same Ti/Al and Zr/Al patterns occur in Miocene sections from the island of Gavdos, south of Crete [Schenau *et al.*, 1999]. These differences were attributed to variations in the input of fluvial and aeolian detritus to these sites, with finer-grained sediments of fluvial derivation being more important during sapropel deposition and coarser aeolian material being much more important during marl deposition.

The distribution of Ba/Al ratios is closely coherent with organic C (Figure 8); most Ba enrichments above crustal abundances ($\text{Ba/Al} = -0.0068$) [Taylor and McLennan, 1985] occur in the sapropels. In some cases, Ba is also enriched just above a sapropel, such as S1 (Si2), S3 (Si8), and S6 (Si16), and

Figure 8. Down core distributions of Sr/Al, Rb/Al, Zr/Al, and Ba/Al. Variations in Sr/Al are coherent with those of Ca/Al and Mg/Al, whereas variations in Rb/Al and Zr/Al follow K/Al and Si/Al, respectively. Ba/Al peaks are confined to each of the sapropels but also occur between S5 and S6, between S9 and S10, and below S6 and S10.

there is a prominent Ba/Al maximum below S10 (Si30). The marls have Ba/Al ratios of roughly 0.003 upon which the marked enrichments in the sapropels are superimposed. Almost all the Ba in the S1 sapropel is present as 1–3 μm barite crystals [Thomson *et al.*, 1995], similar to its mode of occurrence in open ocean pelagic sediments [Church, 1970; Paytan *et al.*, 1993; Ginge and Dahmke, 1994].

Down core variations in total Ba abundances record palaeoproductivity variations [Schmitz, 1987], and palaeoproductivity algorithms using total sedimentary Ba concentrations have been developed [Dymond *et al.*, 1992; François *et al.*, 1995]. Although there has been some minor diagenetic redistribution of Ba in core MD 84641 [van Os *et al.*, 1991], as seen, for example, below S10 (Si30), the close coherence between Ba/Al and organic C suggests higher production during sapropel formation [Thomson *et al.*, 1999]. Note, in addition, that there are small Ba/Al peaks between S9 (Si22) and S10 (Si30) where Si/Al and Zr/Al ratios are low.

Low and more or less constant Mn/Al ratios occur in the marls, whereas sporadic high values occur at the bases and the tops of several of the sapropels, most notably below S6 (Si16), above and below S10 (Si30), and above S1 (Si2) (Figure 9). Small enrichments are also seen in stages 3, 8, and 11. Since manganese in sediments is strongly affected by the redox state of the sediment [Lynn and Bonatti, 1965], the pattern of Mn distribution is controlled almost entirely by postdepositional reaction. Many eastern Mediterranean sapropels have been altered by the

downward diffusion of pore water oxygen after the formation of the sapropels ceased and the marl-forming regime was re-established [de Lange *et al.*, 1989]. The effects of this process are similar to those in organic-rich North Atlantic turbidites [Wilson *et al.*, 1985; Thomson *et al.*, 1993], in that the organic material in the upper part of the sapropel is degraded and redox-sensitive metals and metalloids are redistributed around the downward migrating oxidation front. Sapropels may be completely removed by this process [Higgs *et al.*, 1994]. The details of these effects in the youngest Mediterranean sapropel (S1) have been fully described [Pruysers *et al.*, 1993; Thomson *et al.*, 1995; Van Santvoort *et al.*, 1996].

Several other redox-sensitive trace metals can be redistributed by the processes just described. Thus Cu/Al, Cr/Al, and Zn/Al (Figure 9) have down core variations that are poorly correlated with the positions of the sapropels. Isolated single-point peaks in the ratios are found in the marls, some of which are coincident with minima in Si/Al and Zr/Al (Figures 5 and 8). In the case of Zn/Al, maxima and minima occur within different sapropels. Down core variations in Co/Al, Ni/Al, Mo/Al, and to a lesser extent V/Al (Figure 10) are much more closely related to the sapropel boundaries, maxima most often occurring within these horizons. Some isolated peaks of Co/Al and V/Al are found in the marls, whereas Ni/Al and Mo/Al maxima are largely confined to the sapropels; Ni/Al begins to increase immediately below some of the sapropels, whereas Mo/Al ratios show the best coherence with the boundaries of the sapropels. In addition, Co/Al, Ni/Al,

Figure 9. Down core distributions of Mn/Al, Cu/Al, Cr/Al, and Zn/Al. The lack of a clear relationship between the distribution of these ratios and the occurrence of sapropels suggests that postdepositional processes have largely obliterated any signals of changing sedimentary conditions.

and Mo/Al ratios are much lower in the upper section of the composite sapropels S4 (Si10), S6 (Si16), and S8 (Si20) than they are above the intercalated organic-poor mud layer.

The enrichment of a wide range of trace metals in the sapropels of core MD 84641 that are removed to bottom sediments under reducing conditions suggests that bottom waters or the interstitial environment of the sediment was oxygen-deficient or even sulphidic during their formation. It is not possible to deduce the degree of oxygen deficiency with the elemental data available because it is known that enrichments of many of the same metals can occur by the downward diffusion of a metal from oxic bottom waters into anoxic subsurface horizons [Calvert and Pedersen, 1993]. Other evidence can sometimes be used to infer bottom water anoxia, such as the presence of laminations or the complete absence of benthos [Jorissen, 1999], and both of these indicators can be found in Mediterranean sapropels. However, there are situations where the sapropels contain a benthic foraminiferal fauna, albeit adapted to very low oxygen conditions [Brolsma, 1978; Jonkers, 1984; Oggioni and Zandini, 1987; Vismara Schilling and Coulbourn, 1991], and one must always be cognizant of the fact that fine laminations can be imparted to sediments under oxic conditions by mat-forming diatoms [Kemp and Baldauf, 1993] that have been found in S5 (Si12) [Kemp et al., 1999]. Anoxic bottom water conditions during sapropel formation are also indicated by the sulphur isotopic composition of pyrite in several sapropels [Passier et al., 1999]. In a similar manner to the relatively high bottom oxygen conditions prevailing during the deposition of an upper sapropel

section in the Adriatic Sea inferred from the benthic fauna [Rohling et al., 1997], we also infer nonanoxic conditions during the deposition of the upper sections of sapropels S4 (Si10), S6 (Si16) and S8 (Si20) in core MD 84641 from the very low Co/Al, Ni/Al and Mo/Al ratios and their contrast to the much higher values in the lower sections of the same sapropels.

3.5. Sapropel-Marl Differences

Apart from the organic carbon and nitrogen contents, the mineralogical and chemical composition of the sapropels and marls in core MD 84641 shows that the two facies are compositionally distinct. The differences can be summarized as follows:

1. The marls contain abundant high-Mg calcite, whereas the sapropels have much lower or undetectable amounts of this phase.
2. The sapropels are finer grained than the marls, as shown by their lower quartz/clay, Si/Al, and Zr/Al ratios.
3. The fine-grained (clay) aluminosilicate compositions are different, sapropels containing more smectite and kaolinite and marls containing a higher proportion of illite.
4. The sapropels have high concentrations of trace metals that are precipitated under anoxic conditions as their respective sulphides (Co, Mo, and Ni) or are reduced to their lower oxidation states under low redox conditions and are fixed in the sediment by precipitation of a reduced phase or adsorption onto other solid phases (Mo, V). The distributions of other metals normally found enriched in anoxic conditions (Cr, Cu, and Zn) have been seriously affected by diagenetic redistribution.

Figure 10. Down core distributions of Co/Al, Ni/Al, Mo/Al, and V/Al. In contrast to the distributions shown in Figure 9, Co, Ni, Mo, and V maxima occur in sapropels, apart from Co in S1 and S9 and V between S9 and S10.

5. The sapropels have significantly higher concentrations of Ba compared with the marls.

The differences in the mineralogy and the bulk composition of the terrigenous fraction of the sediments indicate that the source or sources of the sediment in the sapropels and marls were different. Consequently, the sapropel record in core MD 84641 cannot be interpreted simply as the imposition of high organic supply or preservation upon a uniform compositional background. These results are more plausibly interpreted as a change in the conditions of sedimentation and/or sediment delivery, possibly linked to changes in the circulation in the eastern Mediterranean between periods of marl and sapropel formation.

Holocene sediment sources in the eastern Mediterranean are dominated by supply from the Nile and from the southern arid land areas via wind transport [Venkatarathnam and Ryan, 1971; Dominik and Stoffers, 1978]. Thus the fine-grained material in the deep water sediments is dominated by smectite (via the Nile) and kaolinite (from rivers and the deserts via the wind). Illite and chlorite are much more abundant in the western part of the eastern basin, these minerals having a northern source. The higher smectite and kaolinite contents of the sapropels suggest that sediment supply from the southern source was even greater during the deposition of sapropels at the site of core MD 84641, consistent with the higher flow rates of the Nile at these times [Rossignol-Strick *et al.*, 1982; Rossignol-Strick, 1983; 1985]. This has recently been confirmed from the distribution of $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in the surface sediments of the Levantine Basin [Krom *et al.*, 1999a] and from $^{87}\text{Sr}/^{86}\text{Sr}$ and major element ratios in S1 (Si2) and associated marls in cores from the same region [Krom *et al.*, 1999b]. These authors showed that the compositional and isotopic data require a simultaneous decrease in Saharan dust

input and an increase in Nile River input during the deposition of S1 (Si2).

As well as having a different clay mineral composition, the marls between each of the sapropels in core MD 84641 are also coarser grained, as shown by the higher quartz/clay, Si/Al, and Zr/Al ratios. The highest ratios in the entire core are found immediately below each of the sapropels, especially S1 (Si2) and S5 (Si12). It is unlikely that hydrodynamic reworking has caused this contrast in view of the water depth at the core site. Because of the distance from fluvial sources, aeolian transport is the most likely transportation agent. At the present time, winds transport large amounts of dust over the eastern and central Mediterranean from the Sahara [Ganor and Mamane, 1982] and from northern sources [Tomadin *et al.*, 1984], and wind transport has been implicated in the distribution of the fine fraction to the sediments of the eastern basin [Venkatarathnam and Ryan, 1971; Tomadin, 1981; Wehausen and Brumsack, 1999]. Thus a more vigorous glacial atmospheric circulation, as generally deduced for the LGM [Petit, 1981; Mahowald *et al.*, 1999] together with a change in the sedimentary source areas suggest that the two sedimentary facies in core MD 84641 reflect different climatic and oceanographic regimes.

The distinct compositional differences between sapropels and marls in MD 84641 provide a means for identifying the positions of so-called missing sapropels. Cita *et al.* [1977] stated that sapropels S2 (Si4/6) and S4 (Si10) are only "weakly expressed" in cores from the ridge crest of the Ionian Sea and that S9 (Si22) and S10 (Si30) may be very thin or smeared by bioturbation. Rossignol-Strick, [1985] found that, on the basis of the close relationship between the occurrence of sapropels and African climate changes driven by precessionally forced monsoon

variability (to be discussed in section 3.7), sapropels should occur at 292 and 314 kyr in the eastern Mediterranean, but they are absent. Thus it appears that there are genuine gaps in the sapropel record in the eastern Mediterranean. Postdepositional "burn-down" [de Lange *et al.*, 1989] is one process that could remove a sapropel from the section, thin or relatively organic-poor horizons being especially susceptible [Higgs *et al.*, 1994].

It is necessary to distinguish between situations where sapropels did not form because the climatic conditions did not change sufficiently and situations where such deposits did form but have been completely removed by the burn-down mechanism. In the latter case, horizons of former sapropels are clearly marked by finer sediment grain size and distinctive mineralogy, properties that are not altered by postdepositional oxidation. Thus Figures 4, 5, 6, and 8 show that there are prominent minima in quartz/clay, Mg calcite/calcite, Si/Al, Ti/Al, K/Al, Mg/Al, Rb/Al, and Zr/Al ratios, all indicators of sapropel composition, in stages 3, 8, and 9 where there are no organic carbon maxima. The minima that coincide with or lie close to Ba/Al peaks, such as between S9 (Si22) and S10 (Si30) (Figure 8), may represent a residual signal of sapropel formation; these are the "ghost" sapropels of other authors. Other minima, notably between S1 (Si2) and S3 (Si8) (Figures 5 and 8), possibly record periods of altered climatic and oceanographic conditions that were insufficient to trigger sapropel formation. These are genuine missing sapropels. The recognition of a reliable conservative tracer of sapropel formation conditions in core MD 84641 provides further insight into the relationship between sapropel formation and palaeoclimatic changes over the last 460 kyr, which will be discussed in section 3.7.

3.6. Composite Sapropels

In core MD 84641, sapropels S4 (Si10), S6 (Si16), and S8 (Si20) contain an intercalated horizon of organic-poor sediment; they are therefore composite sapropels [Rohling *et al.*, 1997]. Organic C concentrations in S6 (Si16) decrease to levels characteristic of the marls, whereas they are significantly higher than the marl values in the organic-poor layers in S4 (Si10) and S8 (Si20) (Figure 2). The $\delta^{18}\text{O}_{\text{Ruber}}$ increases significantly in the organic-poor layer in S4 (Si10) and S8 (Si20), whereas there is no significant change in S6 (Si16). CaCO_3 contents increase slightly in the organic-poor layer in S8 (Si20) compared with the sapropelic sediments above and below, whereas there is no significant change in S4 (Si10) and S6 (Si16). High-Mg calcite increases notably in S6 (Si16), there is a small increase in S4 (Si10), and this phase could not be detected in S8 (Si20) (Figure 4). Ba/Al ratios closely track the organic C changes in the three composite sapropels, whereas Zr/Al ratios in the organic-poor layer in each of the composite sapropels are slightly higher compared with the sapropelic sediments above and below this horizon (Figure 8).

The two sections of the composite sapropels have different minor element compositions. Thus, while Co/Al, Ni/Al, and Mo/Al ratios all have marked maxima in all sapropels, they remain low in the upper sapropel section of S6 (Si16) and S8 (Si20) (Figure 10); only Mo/Al is much lower in the upper part of S4 (Si10). Since the enrichments of these three metals in the sapropels probably reflect anoxic bottom water and/or interstitial conditions during sapropel formation, the low degrees of enrichment in the upper section of S4 (Si10), S6 (Si16), and S8

(Si20) must be due to somewhat more oxygenated bottom water conditions following the interruption of sapropel deposition. Note that Ba/Al ratios are higher than background in both sections of the composite sapropels (Figure 8), suggesting a return to higher-productivity conditions after sapropel formation was interrupted by better bottom water ventilation.

Rohling *et al.* [1997] described a similar organic-poor horizon in sapropel S1 (Si2) from the Adriatic Sea, and De Rijk *et al.* [1999] found a possibly coeval layer in the southern Aegean Sea. These authors interpreted high-resolution benthic and planktonic foraminiferal data as reflecting a small sea surface cooling and increased bottom water ventilation that interrupted sapropel deposition at this time. Their data also showed that there was a significant increase in organic carbon flux to the seafloor (ecosystem enrichment) concomitantly with a progressive decrease in bottom water oxygenation for the lower section and more oxygenated bottom water conditions during the deposition of the upper section of the sapropel.

The combined data from MD 84641 suggest that productivity decreased (Ba/Al decreases), salinity increased (Mg calcite/calcite increases), wind speed increased (Zr/Al increases), and bottom water and/or interstitial oxygenation increased (Co/Al, Mo/Al, and Ni/Al decrease) during the interruption to sapropel S4 (Si10), S6 (Si16), and S8 (Si20) formation. Moreover, bottom water oxygenation remained high during the deposition of the upper sections of these composite horizons even though production returned to the high levels typical of the other noncomposite sapropels in MD 84641. Subtle changes in the hydrological balance during periods when sapropel formation was otherwise favored were evidently responsible for the formation of a number of composite sapropels.

3.7. Climatic Control of Sapropel Formation

The formation of the eastern Mediterranean sapropels has been explained by several different models of climatic and/or oceanographic change, and these have been recently reviewed [Rohling and Hilgen, 1991; Rohling, 1994]. All explanations involve a drastic change in the hydrological balance relative to the modern situation, which is a simple antiestuarine flow of surface Atlantic inflow balanced by evaporative loss and deep water return flow through the Gibraltar Straits [Wüst, 1961; Lacombe and Tchernia, 1972]. Olausson [1961] proposed that the influx of fresh water from the surrounding land and especially from the Black Sea, following its postglacial reconnection with the Mediterranean through the Bosphorus and Dardanelles Straits, led to a decrease in the surface salinity in the eastern basin of the Mediterranean, and this caused a density stratification sufficient to impede or halt bottom water formation. Kullenberg [1952] had earlier argued, by analogy with the stagnant basins of the modern Baltic, that both the supply of oxygen to the bottom waters should be low and the supply of organic matter should be high for organic- and sulphide-rich layers to form in the sediments of the eastern Mediterranean. Thus the lack of replenishment of dissolved oxygen to the deep basin waters due to water column stratification has become the widely accepted model for sapropel formation.

Temperature-corrected oxygen isotope anomalies of planktonic foraminifera have been interpreted as showing that sea surface salinities (SSS) were significantly lower during sapropel formation [Williams *et al.*, 1978], a potential cause of the larger

range of Mediterranean foraminiferal $\delta^{18}\text{O}$ values than is found in open ocean cores [Vergnaud-Grazzini *et al.*, 1986]. More recent analyses of the isotopic foraminiferal record [Kallel *et al.*, 1997b; Rohling and De Rijk, 1999] support the interpretation of significantly lower SSS during the formation of S1 (Si2). Kallel *et al.* [1997b] concluded that SSS was more or less uniform throughout the eastern and western basins and that surface density was higher in the west, leading to inflow of deep water from the western basin. In contrast, Rohling and De Rijk [1999] found that an east-west salinity gradient is still consistent with the data, although there appears to have been a marked water $\delta^{18}\text{O}$ depletion in the Levantine Basin most likely due to the influence of high Nile discharge [Rossignol-Strick *et al.*, 1982]. Since water $\delta^{18}\text{O}$ is not a conservative property controlled solely by the precipitation-evaporation balance, Rohling and De Rijk, [1999] were reluctant to provide estimates of the latter for S1 (Si2) time because the translation of water $\delta^{18}\text{O}$ changes to sea surface salinity changes is not straightforward. Finally, the modeling results of Myers *et al.* [1998] suggest that the adoption of the inferred S1 (Si2) salinity changes of Thunell and Williams [1989], Kallel *et al.* [1997a] and Rohling and De Rijk [1999] leads to solutions that have antiestuarine flow, as at the present, throughout the Mediterranean.

Rossignol-Strick [1985] and Rossignol-Strick *et al.* [1982] correlated the temporal record of sapropels to large increases in freshwater input from the Nile River, which respond directly to monsoonal rainfall over eastern tropical Africa. Rossignol-Strick [1987] also suggested that increased runoff from Eurasian rivers contributed to the change in the hydrological balance in the eastern Mediterranean, and this is supported by the oxygen

isotopic composition of Late Pleistocene [Jenkins and Williams, 1984] and Pliocene [Gudjonsson and van der Zwaan, 1985] planktonic foraminifera in the sapropels. Evidence for much more humid conditions in the eastern Mediterranean during S1 (Si2) time also comes from pollen records from North Africa [Street-Perrott and Perrott, 1993] and southern Europe [Wijmstra *et al.*, 1990; Cheddadi *et al.*, 1991; Rossignol-Strick, 1995] as well as records of North African and Levantine lake levels (summarized by Yu and Harrison [1996]). Bar-Matthews *et al.* [1999] have reviewed existing knowledge on the wetter climate of the Negev Desert during the last 140 kyr based on the isotopic composition of speleothems. These data are in good agreement with the results of climate model simulations of Holocene climates over Africa [Kutzbach and Otto-Bliesner, 1982; Kutzbach and Guetter, 1986]. During the formation of S1 (Si2) between 7.3 and 9.3 kyr ago, increased solar radiation at 30°N , due to the occurrence of perihelion in July and an axial tilt of 24.2° (compared to the modern value of 23.4°), led to a considerable strengthening of the land-ocean pressure gradient. This caused stronger onshore air flow into northern and eastern Africa, which led to increased humidity and a larger effective moisture (precipitation minus evaporation) flux over the land [Kutzbach and Street-Perrott, 1985]. Thus monsoonal rainfall over the Ethiopian Highlands was much greater, leading to greater runoff into the eastern Mediterranean [Rossignol-Strick, 1983]. From older paleolake records this pattern of increasing humidity in the Sahara/Sahel region appears to have been repeated in stages 5, 7, and 9 [Szabo *et al.*, 1995], coincident with the formation of sapropels in core MD 84641 at these times. Sapropel formation in the eastern Mediterranean is also coeval with marked decreases in SSS in the

Figure 11. Time scale of sapropel occurrence in core MD 84641 and summer insolation at 65°N . Insolation data (W/m^2) derived from the Laskar90 series [Laskar, 1990] by means of the AnalySeries software [Paillard *et al.*, 1996]. Observable sapropels, as defined by the organic C maxima, are indicated by the stippled pattern, and the positions of the "ghost" and missing sapropels, as defined by minima in Si/Al, are indicated by the oblique line pattern. The insolation-based sapropel numbering scheme of Langeris *et al.* [1997], even numbers corresponding to insolation maxima, is indicated on the left, and shows that all events that could have led to sapropel formation are present in core MD 84641.

Andaman Sea due to increased monsoonally driven runoff from the Indian subcontinent [Kallel *et al.*, 2000], testifying to the enhanced atmospheric transport of moisture from ocean to land.

According to Thunell and Williams [1989], the SSS contrast between sapropel S1 (Si2) and the LGM amounts to 2-3‰. The inferred higher LGM SSS is consistent with extensive information on the existence of dry and cold open steppe-like conditions over large tracts of North Africa at the LGM [Street-Perrott and Perrott, 1993]. The most recent modeling study [Rohling, 1999] has found that average glacial salinities were probably 41-42‰ and, by analogy with the modern situation, somewhat higher in the eastern basin, consistent with the presence of abundance high-Mg calcite in eastern Mediterranean marls. High-Mg calcite also occurs abundantly in Red Sea sediments [Gevirtz and Friedman, 1966] where current bottom water salinities are ~40 ‰ [Locke and Thunell, 1988] owing to high evaporation rates. Although the inferred higher LGM SSS does not necessarily mean that the climate was more arid (because of the complicated relationship between evaporation rate and water exchange), the indication of greater aeolian input (higher quartz/clay, Si/Al, and Zr/Al) during these intervals in core MD 84641 does point to somewhat drier conditions on land or higher wind speeds.

The timing of sapropel formation with respect to the relevant insolation timeseries is shown in Figure 11. Maxima in summer insolation correspond to maxima in the monsoon index [Rossignol-Strick, 1983, 1984], the insolation gradient between the equator and the tropics, which drives more intense monsoonal circulation. Organic carbon peaks in each of the sapropels lag the insolation maxima by 3-9 kyr. Hilgen [1991] found that there was a 4 kyr lag between the sapropels S1 (Si2) to S12 (Si44) and the peaks of the precession index of Berger [1978], whereas Lourens *et al.* [1996] introduced a 3 kyr lag between the midpoints of sapropels and summer insolation maxima of Laskar [1990]. The significant and reproducible lag of sapropel formation evidently reflects a lag in the thermal contrast between the tropics and the equator driven by an offset in summer solstice-perihelion alignment [Short and Mengel, 1986]. These authors also pointed out that the peak thermal contrast 3 kyr after maximum insolation translates into a greater moisture availability in the rainy season, amounting to a 50% change in saturation vapor pressure, pointing to a significant impact of orbital variations on the tropical hydrological cycle.

Variations in Si/Al in core MD 84641 are coherent with changes in summer insolation at 65°N (Figure 11). As discussed previously, minima in the ratio mark the positions of each of the sapropels in this core. However, in addition, there are clear minima in stages 8 and 9, where sapropels are absent, that are lagged by 2-3 kyr with respect to insolation and very small, but significant, minima in stage 3, where there is a low-amplitude double insolation minimum. Thus the Si/Al ratio appears to be well tuned to the insolation curve and provides a secure record of conditions that could have led to the formation of sapropels in the eastern Mediterranean. This provides a more complete record of sapropel-forming conditions than the down core presence/absence of organic-rich sapropels.

3.8. Glacial Sapropels

The apparently anomalous occurrence of sapropel S6 (Si16) in glacial stage 6 has been pointed out by Cita *et al.* [1977] and

Rossignol-Strick [1983]. The $\delta^{18}\text{O}_{\text{ruber}}$ data from core MD 84641 (Figure 2) as well as several other cores from the eastern Mediterranean clearly show that this stage was fully glacial. Moreover, the abundance of *G. ruber* decreases drastically and *G. bulloides* increases at this horizon, consistent with much cooler surface water conditions [Kallel *et al.*, 2000]. As discussed in section 3.7, the trigger for sapropel formation is evidently a drastic change in the hydrological balance of the eastern Mediterranean, driven by precessionally modulated insolation changes. Evidence of such changes during glacial as well as interglacial stages is not difficult to find in view of the timing of the major glacial-interglacial fluctuations and the insolation record over the last 800 kyr. Low-latitude (23.5°) summer insolation during stage 6 was as high as that during events 5.1 and 5.3 and only slightly lower than that during events 5.5 and 7.1 [Mélières *et al.*, 1997]. Thus, in spite of a fully glacial climate condition at high latitudes the strong insolation maximum at 175 kyr was sufficient to change the hydrological state of the eastern Mediterranean and induce the formation of a sapropel (S6 or Si16) at ~168 kyr. The driving force for such hydrological changes is in the low latitudes (the monsoons and subtropical precipitation patterns), and these are controlled mainly by precessional insolation changes [Rossignol-Strick, 1985]. Possible reasons for the fact that the climate remained in a glacial state in spite of the relatively high insolation maximum at 175 kyr have recently been addressed by Ridgwell *et al.* [1999].

3.9. Palaeoproduction During Sapropel Formation

The geochemical data can be used to estimate plankton production during the sapropel events in the eastern Mediterranean. The sedimentary accumulation rate of excess Ba (relative to the aluminosilicate background) in S1 (Si2), using the ^{14}C -derived linear sedimentation rate of S1 (Si2) (data from Fontugne *et al.* [1994]) and the concentration of biogenous Ba (the excess relative to a Ba/Al ratio of 0.003), implies a palaeoproduction of $150 \text{ g C/m}^2 \text{ yr}^{-1}$ using the barite-based algorithm of Paytan *et al.* [1996]. Using the method of Dymond *et al.* [1992] to derive the water column settling flux of biobarium (which takes account of the degree of preservation of Ba in the sediment) and the relationship between biobarium flux and new production (which can be equated with the export flux of organic matter below the photic zone [Eppley and Peterson, 1979]) defined by François *et al.* [1995] gives an estimate of new production of around $5.3 \text{ g C/m}^2 \text{ yr}^{-1}$. These two values suggest that production and organic settling fluxes during Si (Si2) were roughly an order of magnitude larger than modern values from the central eastern basin, which are of order $10\text{-}20 \text{ g C/m}^2 \text{ yr}^{-1}$ and $<1 \text{ g C/m}^2 \text{ yr}^{-1}$, respectively [Berman *et al.*, 1984a, 1984b]. It is unlikely that the very low production levels determined today in the eastern basin are not representative of the production conditions prior to the completion of the Aswan High Dam in 1965 because the high nutrient levels and high phytoplankton cell counts off Egypt before 1965 were confined to a relatively narrow strip <60 km wide off the river mouth, the offshore region being an "unproductive" zone [Halim *et al.*, 1967].

We have not attempted to reconstruct palaeoproduction levels from the accumulate rate of organic carbon in the sediment [Müller and Suess, 1979; Sarnthein *et al.*, 1987] because the carbon accumulation rate in marine sediments is tightly controlled by the linear sedimentation rate [Middelburg *et al.*, 1997]. The

calibration of the method for determining palaeoproduction from sedimentary carbon accumulation rates therefore produces results that largely reflect sedimentation rate variations rather than the settling flux of organic matter from the sea surface.

The higher production during sapropel formation that we have deduced from the geochemical evidence is strongly supported by the results of studies of planktonic foraminifera [Rohling, 1989; Lourens et al., 1992], ostracodes [van Harten, 1987], calcareous nanofossils [Castradori, 1993], and diatoms [Schrader and Matherne, 1981; Kemp et al., 1999]. The latter authors speculate that diatom fluxes could well have been high during all sapropel periods in spite of the poor preservation of diatom opal in the sediments because of the low dissolved silicon concentrations in deep Mediterranean waters. The diatom groups preserved (rhizosolenids and *Hemiaulis*) are typical of stratified waters where high production is maintained toward the base of a deep photic zone. The results of the modeling study of Stratford et al. [2000] suggest that a full circulation reversal of the eastern Mediterranean [Berger, 1976] is required to increase production sufficiently for sapropels to form over the entire basin.

4. Conclusions

Mediterranean sapropels are organic-rich, fine-grained muds that formed during periods of greatly altered freshwater balance. The mineralogical and chemical compositions of these deposits are significantly different from the associated marls; they are finer grained, and they contain diagnostic clay mineral phases that point to a southern source or sources. Marls lying immediately below the sapropels are coarser grained and they contain high concentrations of high-Mg calcite together with clay minerals characteristic of northern and/or western sources. These contrasts

suggest that transport agencies (most likely aeolian) for the terrigenous sediment fraction were more vigorous during the accumulation of the marls and that deep water salinities were quite different at times of sapropel and marl deposition. These features can be reconciled if conditions in the eastern Mediterranean were more arid during marl formation, leading to more evaporative water loss and higher wind speeds. Primary production was higher during sapropel formation, causing a large increase in the settling flux of organic matter to the basin floor. This would have drawn down deep water oxygen levels, which also might have been replenished less frequently if circulation restricted deep water renewal. The sediments themselves were anoxic during sapropel formation, although the upper sections of composite sapropels show clear evidence of oxygenated bottom waters even though production was high. The positions of missing and ghost sapropels can be identified from the down core distributions of lithogenous tracers (quartz/clay, Si/Al, and Zr/Al ratios) that are not affected by postdepositional diagenetic modification of the sedimentary record. The reconstructed record shows that the conditions for potential sapropel formation occurred slightly after every significant insolation maximum during the last 350 kyr.

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