20. PLIOCENE AND PLEISTOCENE ABUNDANCE AND PRESERVATION OF SILICEOUS MICRO-FOSSIL ASSEMBLAGES FROM SITES 794, 795, AND 797: IMPLICATIONS FOR CIRCULATION AND PRODUCTIVITY IN THE JAPAN SEA¹

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ABSTRACT

A close examination of the siliceous microfossil assemblages from the sediments of ODP Leg 127, Japan Sea Sites 794, 795, and 797, reveals that upper Pliocene and Pleistocene assemblages have been subjected to more dissolution than have lower Pliocene assemblages. This conclusion is based on semiquantitative observations of samples processed for diatoms and radiolarians. Although preservation of opaline microfossils in some upper Pliocene and Pleistocene samples is better than others, in general, the poorly preserved state of these assemblages supports the notion that opal dissolution, in response to lowered productivity, is responsible for the paucity of siliceous microfossils in upper Pliocene and Pleistocene sediments. The lithological transition from diatomaceous oozes to silts and clays corresponds to a change between dominantly well preserved more poorly preserved siliceous assemblages, and is termed the late Pliocene Japan Sea opal dissolution transition zone (ODTZ). The base of the ODTZ is defined as the uppermost occurrence of high abundances of moderately to well preserved valves of the diatom *Coscinodiscus marginatus*. The dissolution transition zone is characterized by partially dissolved refractory assemblages of radiolarians, the presence of *C. marginatus* girdles, *C. marginatus* fragments, siliceous sponge spicules, and a general decrease in weakly silicified, less solution resistant diatoms upward in the section. The top of the dissolution transition zone marks the level where whole *C. marginatus* valves and *C. marginatus* fragments are no longer present in significant numbers.

Dissolution of the late Pliocene and Pleistocene opaline assemblages is attributed mainly to changes in paleoceanographic circulation patterns and decreased nutrient (dissolved silicon) contents of the water column, and possibly dissolution at the sediment/water interface, rather than to post-depositional dissolution or diagenesis. We suggest that the transition from silica-rich to silica-poor conditions in the Japan Sea was due to fluctuations of deep-water exchange with the Pacific through the Tsugaru Strait between 2.9 and 2.3 Ma.

INTRODUCTION

A primary purpose of Leg 127 was to determine the paleoceanographic evolution of the Japan Sea by examining the record of microfossil assemblages, in particular siliceous microfossils. A conspicuous feature of Japan Sea sediments is the diagenetic transformation of opal-A to opal-CT between 200 and 300 mbsf, caused by optimal conditions of sediment thickness, age, and heat flow (Langseth and Tamaki, this volume). Unfortunately, the relatively shallow occurrence of the opal-A/CT transition has obliterated most of the paleontological record of siliceous sedimentation below the opal-A/CT boundary, limiting the age of preserved opaline microfossils in the sediments cored during Leg 127 to no older than late Miocene or ~8 Ma.

In general, above the opal-A/opal-CT diagenetic boundary the upper Miocene and lower Pliocene sediments of all four sites (Fig. 1) sampled in the Japan Sea during Leg 127 are rich in siliceous microfossil remains (diatom silty clays and oozes). The siliceous microfossils found in these sediments (diatoms, radiolarians, and siliceous sponge spicules) are moderately to well preserved and are typically common to abundant. In contrast, upper Pliocene and Pleistocene sediments of the sites contain low percentages of diatoms and are composed mainly of silts, clays, and ashes. Furthermore, uppermost Pleistocene sediments display distinctive alternating light and dark layers, with siliceous microfossils showing variable states of abundance and preservation.

PURPOSE OF STUDY

Because of the stark contrast between relatively silica-poor upper Pliocene and Pleistocene sediments and silica-rich lower Pliocene and upper Miocene sediments, and the assumption that high siliceous plankton productivity in the water column is accompanied by high opal contents in the underlying sediments (Leinen et al., 1986), the Shipboard Scientific Party (1990b-e), and Tamaki, Pisciotto, Allan, et al. (1990) concluded that the Miocene and early Pliocene waters of the Japan Sea were very productive. Preliminary surveys of samples from the diatomaceous oozes revealed large abundances of the diatom Coscinodiscus marginatus. The presence of this diatom complicated assemblage descriptions because it is not present in such high proportions in plankton samples in the North Pacific (Sancetta, 1982), suggesting it is a dissolution-resistant form (Burckle, Sturz, and Emanuele, this volume). Our approach in this paper is to describe and interpret the differences in the siliceous microfossil assemblages over the interval between the diatom-rich oozes and the diatom-poor clays and silts in order to determine the nature of Pliocene and Pleistocene paleoceanographic change in the Japan Sea.

The Record of Coscinodiscus marginatus

Coscinodiscus marginatus is a conspicuous component of the sediments of Japan Sea Sites 794, 795, and 797. This diatom is abundant (~50% of the diatom assemblages) in the clayey diatomaceous oozes. Because it is quite large ($30-100 \mu m$ in diameter), it comprises a large proportion of the >63 µm size fraction. Coscinodiscus marginatus is both a paleoenvironmental indicator and a dissolution indicator (Schrader, 1973) as well as an important component in North Pacific assemblages (Sancetta and Silvestri, 1986). Coscinodiscus marginatus is considered to be transitional between subtropical and subarctic water masses (Sancetta and Silvestri, 1986). Sancetta and Silvestri (1986) note its abundance in a piston core of calcareous-siliceous ooze from the northeastern Pacific (Core V20-119) between

¹ Pisciotto, K. A., Ingle, J. C., Jr., von Breymann, M. T., Barron, J., et al., 1992. Proc. ODP, Sci. Results, 127/128, Pt. 1: College Station, TX (Ocean Drilling Program). ² Department of Cascinger See Ference and Cascing Program.

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Figure 1. Map of the Japan Sea showing the location of ODP Leg 127 Sites 794-797.

~2.6 and 2.4 Ma and suggest it was deposited during times of high productivity in the surface waters. Furthermore, Sancetta and Silvestri (1986) note the decline of *C. marginatus* after 2.4 Ma to minimum values at ~2.0 Ma. The investigators conclude that strong yet fluctuating degrees of stratification of North Pacific surface water masses occurred at this time as a result of glacial-interglacial cycles. Our observations suggest that the overall abundance and preservation of diatoms and radiolarians decreases following peak abundances of *C. marginatus*, and generally corresponds to the lithologic break between diatom oozes and diatom clays. Furthermore, the presence of dissolution-resistant radiolarians such as Trissocyclids (similar to what was observed by Holdsworth and Harker, 1975) in the Pleistocene diatom clays with variably abundant and fragmented *C. marginatus* frustules, is in agreement with increased dissolution at this time.

Opal Dissolution Transition Zone (ODTZ)

The opal dissolution transition zone (ODTZ) is an interval we recognize in the Japan Sea cores that marks the change between dominantly well preserved to more poorly preserved siliceous assemblages. It generally corresponds to a lithological change between diatomaceous oozes and diatom silts and clays. We define the base of the ODTZ as the uppermost occurrence of high abundances of whole *C. marginatus* valves or "*Coscinodiscus* oozes" (Burckle, this volume). Within the ODTZ the following components are common: partially dissolved and broken *C. marginatus* valves and girdles; siliceous sponge spicules; and partially dissolved radiolarians. The top of the ODTZ is marked by the decrease in the numbers of whole *C. marginatus* and *C. marginatus* fragments. We discuss the nature of the ODTZ at each site and its paleoceanographic significance in the results and discussion sections.

OCEANOGRAPHIC SETTING

The Japan Sea is a relatively young back-arc basin that began opening in the late Oligocene or early Miocene (Leg 127 and Leg 128 Shipboard Scientific Parties, 1990). Since this time, complex tectonic activity has resulted in changes in the shape and boundaries of the basins of the Japan Sea (Iijima and Tada, 1990), which have undoubtedly affected the paleoceanography and depositional conditions of this marginal sea. The Japan Sea is divided into two deep basins: the Japan Basin to the north (maximum water depth ~3700 m, Tamaki, 1988); and the Yamato Basin to the south (Fig. 1, maximum water depth ~3000 m, Tamaki, 1988). The waters in these basins are topographically isolated from Pacific intermediate and deep water masses at the sills of shallow straits. The straits include (from south to north) the Tsushima Strait between Honshu and Korea at a depth of 130 m, the Tsugaru Strait between Honshu and Hokkaido at 130 m, the Soya Strait between Hokkaido and Sakhalin at 55 m, and the Tatar Strait between Sakhalin and Asia at 12 m (Ingle, 1975).

The present day oceanographic and climatic conditions of the Japan Sea keep the waters of the Japan Sea well mixed and well oxygenated (hyperventilated). In the northern region, cold waters with high oxygen content flow south in the Liman Current, sink rapidly, and serve to fill, mix, and ventilate the highly productive basin. Surface waters enter the Japan Sea through the Tsushima Strait as the Tsushima Current (an arm of the warm Kuroshio), and flow out from the Tsugaru Strait (Ingle, 1975; Matoba 1984). Assuming the straits were in approximately the same tectonic positions during glacial times when sea level was lower, the depths of the sills of these straits were shallower and flow was reduced or completely restricted. Studies of latest Pleistocene and Holocene Japan Sea sediments suggest that during certain periods, the deep basins have been stagnant, flow has been cut off at the Tsushima Strait, and water of the Oyashio Current flowed into the basin (see Kobayashi, 1985; Koizumi, 1989; Oba et al., 1991). Such changes in circulation are closely tied to Pleistocene Northern Hemisphere glaciations and may have caused reversals between oxic, anoxic, or suboxic states, and have undoubtedly affected the depositional conditions and sedimentary deposits in the basins of the Japan Sea (Tada and Iijima, this volume).

Four sites were cored in the Japan Sea during Ocean Drilling Program Leg 127. Sites 794, 795, and 796 are located in approximate water depths of 2810, 3300, and 2600 m, respectively, along the eastern margin of the Japan Basin, and Site 797, at a water depth of 2860 m is located in the southwestern part of the Yamato Basin near the Yamato Rise (Tamaki, Pisciotto, Allan, et al., 1990). Today, Sites 797 and 794 are both located beneath the path of the Tsushima current, while Sites 796 and 795, are influenced by the colder more northern waters of the Liman and North Korean Current. Thick sedimentary sections (~500 m thick) were cored at each of these sites with recoveries in unlithified sediments averaging ~75%. Biostratigraphic data suggest the sediments recovered at each site provide a continuous record of Pliocene and Pleistocene biogenic sedimentation.

METHODS

One sample/section was obtained from Holes 794A, 795A, and 797B, down to the opal-A/opal-CT transition, for diatom and radiolarian analysis. Site 796 was not sampled because it is located in an active tectonic zone. The frequent coarse clastic deposits and slumped units which comprise the record at this site make environmental interpretations of the microfossil assemblages suspect (Shipboard Scientific Party, 1990d). The data presented here are restricted to Pliocene and Pleistocene cores where the transition from silica-rich to silica-poor sediments can best be observed.

Diatoms

For this study, samples processed for diatoms were disaggregated in water and a solution of 10% H₂O₂ over low heat for ~10–30 min. The solution was then removed from the heat source and allowed to settle. After sufficient cooling, the liquid was decanted and additional water was added to resuspend the residue. Material was then pipetted from the top 1 mL of suspended residue and strewn onto a cover slip. This randomly settled material was then mounted to a glass slide with Hyrax.

Two slides/sample were made and examined at a magnification of 500× by traversing the long dimension of the slide (30 mm) between 5 and 10 times, depending on the abundance of diatoms in the sample. Positive identification of diatom species was confirmed at a magnification of 1250×. Relative abundances of diatoms reflect percentage estimates of diatom valves relative to other sedimentary material on the slide (Shipboard Scientific Party, 1990a). Similarly, species abundances reflect percentage estimates of the species listed in Tables 1–3 relative to other diatoms in the assemblage. The following convention was used: A = Abundant, >50%; C = Common, 25%–50%; F = Few, 1%–25%; R = Rare, <1%; B = Barren, no diatoms present.

Preservation of diatoms is based on the presence of complete or broken thinly silicified or heavily silicified valves (Shipboard Scientific Party, 1990a). The following three categories were classified:

G = Good: well preserved with numerous delicate diatom frustules M = Moderate: mostly heavily silicified frustules with some thinly silicified valves present

P = Poor: only heavily silicified valves present.

Listed in Tables 1–3 are the abundances of diatoms that are generally diagnostic of Pliocene and Quaternary assemblages of the North Pacific (Koizumi, 1986; Sancetta and Silvestri, 1986). The abundance of *Coscinodiscus marginatus* and *C. marginatus* girdles are listed next to abundance and preservation data on the tables to emphasize how closely fluctuations in the abundance of *C. marginatus* reflect the overall abundance and preservation of the diatom assemblages. Also shown in the tables is the abundance of siliceous

sponge spicules in these slides, which may be a function of increased transport of sediments and/or preservation.

Radiolarians

The collection of quantitative data from the radiolarian assemblages contained in Leg 127 Japan Sea sediments is complicated by large apparent variations in preservation, uncataloged diversity, and taxonomic uncertainties. Therefore, estimates of productivity from these assemblages are tentative, at best. For this study, samples processed for radiolarian paleontology (one sample/section, the >63 µm size fraction of organic and carbonate-free material), were scanned with a light microscope, and estimates of radiolarian preservation and abundance were made on the late Pliocene and Pleistocene intervals (Alexandrovich, this volume). Because Coscinodiscus marginatus is a relatively large diatom (Sancetta, 1982), samples prepared for radiolarian slides (randomly settled refractory material treated with hydrogen peroxide, hydrochloric acid, and washed through a 63 µm sieve), readily recorded the occurrence of this diatom. The abundance of this diatom is so high in the >63 um size fraction of the diatom oozes and clays, the material on the slides becomes several layers thick, which makes the slides look cloudy, and gives them a slightly reddish brown color. These properties were used to distinguish the top of the C. marginatus oozes without microscopic examination, which was then confirmed by microscopic examination.

Above the *Coscinodiscus marginatus* oozes the preservation and abundance of radiolarians decreases, and variations in the abundance of three states (complete frustules, fragmented frustules, and girdles) of *Coscinodiscus marginatus* were observed. Working upsection, a level was noted where *Coscinodiscus marginatus* remains were no longer found. Radiolarians in samples from above this level displayed large variations in assemblage composition, and variable preservation. More than 50% of the samples found above the last significant occurrence of *C. marginatus* were entirely barren of radiolarians. In samples with preserved radiolarians, siliceous sponge spicules were a major component at times.

A transition zone between well preserved late Pliocene and poorly preserved late Pliocene and Pleistocene radiolarian samples was noted in Holes 794A, 795A, and 797B. This zone was also evident in the diatom samples, and observations of both diatoms and radiolarians were used to precisely locate its boundaries (Table 4). Radiolarian samples from this interval were then rescanned, and semiquantitative estimates of the relative contribution of the major biogenic silica components (%radiolarians, %siliceous sponges, %whole *Coscinodiscus marginatus* tests, %fragmented *C. marginatus* tests, and *C. marginatus* girdles) were tabulated (Fig. 4). The presence of terrigenous material such as quartz was also noted in order to determine if samples were subjected to downslope transport.

Chronology

In order to discuss the progression of events observed in the diatom and radiolarian assemblages examined in this study, sample depths were assigned ages using age vs. depth data based on paleomagnetism and (largely diatom) biostratigraphy (Table 5). At the present time, this is the best age control available. Hole 794A has the most complete paleomagnetic record for the Quaternary and Pliocene. Because samples from Sites 794, 795, and 797 contain microfossil assemblages that display high variability in preservation, abundance, and species composition (both siliceous and calcareous), biostratigraphies are not easily tied to standard biozonations and well dated biostratigraphic sequences. In addition, biostratigraphic events (first and last occurrences) tied to the paleomagnetic time scale in the Pacific may not have occurred at the same time in the Japan Sea. Therefore, the age models for the Pliocene and Pleistocene of Sites 795 and 797 may be subject to revision. Until improved age models become available for Leg 127 sites, these age models (Table 5, Fig. 2) are used.

Lithologic unit	Diatom zone	Hole 794A Core, section, interval (cm)	Depth (mbsf)	Abundance	Preservation	Coscinodiscus marginatus	C. marginatus (girdles)	Actinocyclus sp.	Actinoptychus senarius	Azpeitia nodulifer	Chaetoceros spore	Coscinodiscus radiatus	Diploneis bombus	D. smithii	Hemidiscus cuneiformis	Neodenticula kamtschatica	N. koizumii	N. seminae	Nitzschia sp.	Rouria sp.	Rhizosolenia barboi	R. curvirostris	Stephanopyxis turris
IB	N. koizumii N. koi–N. kamt.	8H-1, 124-125 8H-2, 124-125 8H-3, 124-125 8H-4, 124-125 9H-1, 124-125 9H-1, 124-125 9H-2, 124-125 9H-2, 124-125 9H-4, 124-125 9H-5, 124-125 9H-0, 124-125	65.04 66.54 68.04 69.54 72.54 74.54 76.04 77.54 79.04 80.54 82.04	B B C F C A B C C C C A	M M M M M M	R F F F C C C C C	R R R R R R C C C	R	R R R		R R F F	R R F F R F	R R R R	R R R		P	F R F F F F R C P		P		F R F P		P
IIA	Thaslassiosira oestrupii	10H-1, 124-125 10H-2, 124-125 10H-3, 124-125 10H-4, 124-125 10H-6, 124-125 10H-6, 124-125 11H-1, 124-125 11H-2, 124-125 11H-5, 124-125 11H-6, 124-125 12H-1, 123-125 12H-2, 124-125 12H-2, 124-125 12H-3, 124-125 12H-4, 107-108 12H-5, 124-126	84.04 85.54 87.04 88.54 90.04 91.54 93.54 95.04 99.54 101 103 104.5 106 107.4 109 5	A F A A C A C A A A C C C A A	M M G M G M M G M M M M M M M M M M M M	C F C A F F F C A C F C C C	C F A C F F F C C F F R C C	R	R R R		R F R R R F R F F	F	F R R	R F R	R F R	R F F F F F C F	R		F	R	R F R F R F R F R R R R R R		R R R R R R R

Table 1. Abundance and preservation of Pliocene and Pleistocene diagnostic diatoms from Leg 127, Hole 794A, Japan Sea. Also shown is the abundance of sponge spicules throughout the core. Biostratigraphic zonation of the core is after Shipboard Scientific Party (1990b).

Abundances are abbreviated as follows: A = abundant, C = common, F = few, R = rare, and B = barren. For preservation, G = good, M = moderate, and P = poor.

RESULTS

Observations of the siliceous microfossil components in Japan Sea sediments suggest that the late Pliocene lithological change from clayey siliceous oozes to silty clays is accompanied by a transition from well preserved, relatively undissolved biogenic siliceous assemblages, to more poorly preserved, highly dissolved assemblages. The changes in diatoms and radiolarians that accompany this transition are discussed in the following section.

Diatoms

Hole 794A

In Hole 794A, the Pliocene diatoms in Samples 127-794A-12H-6, 124–126 cm (110.5 mbsf), through -8H-3, 124–125 cm (68.04 mbsf), are mostly common to abundant and show moderate preservation (Table 1, Pls. 1 and 2). The assemblages, which are part of the *Thalassiosira oestrupii* Zone to *Neodenticula koizumii–N. kamtschatica* Zone show few to abundant *C. marginatus*, *N. kamtschatica*, *N. koizumii*, *Rhizosolenia barboi*, *Thalassionema nitzschioides*, *Thalassiothrix longissima*, *Thalassiosira oestrupii*, and *T. zabelinae* (Table 1). Upper Pliocene Samples 127-794A-8H-2, 124–125 cm, and -8H-1, 124–125 cm, are barren of diatoms but have rare siliceous sponge spicules.

In the samples processed and examined for diatoms at Hole 794A (<63 µm) the base of the ODTZ, as marked by the uppermost peak of *C. marginatus* abundances, occurs at 88.54 mbsf in Sample 127-794A-10H-4, 124–125 cm (Tables 1 and 4, Fig. 3). The composition

of the sediment at this depth (Unit IB) is silty clay to diatom silty clay (Shipboard Scientific Party, 1990b). Above this depth, *C. marginatus* gradually decreases and their frustules show greater fragmentation (Pls. 2 and 3). In addition, *C. marginatus* girdle bands show the greatest abundance just above the last peak of *C. marginatus* in Sample 127-794A-10H-3, 124–125 cm, at a depth of 87.04 mbsf (Table 1 and Fig. 3). Other trends include increased occurrences of siliceous sponge spicules (few as compared to rare) above the base of the ODTZ (Table 1).

The top of the ODTZ in Hole 794A, as determined from slides examined for diatoms, is in Sample 127-794A-9H-3, 124–125 cm, at a depth of 77.54 mbsf, which is the last common occurrence of *C. marginatus* (Table 1, Fig. 3). Above the top of the transition zone the diversity of diatoms decreases to mostly dissolved frustules of *Coscinodiscus radiatus*, *N. koizumii*, and fragments of *T. nitzschioides* (Table 1), and in addition, the fraction of silt increases.

Hole 795A

In Hole 795A, upper Pliocene and Pleistocene sediments also show decreasing abundances and preservation of diatoms above the maximum occurrence of *C. marginatus* in the diatom silty clays of lithologic Unit IB. Below this horizon, diatom assemblages within the upper Pliocene *T. oestrupii* to *Neodenticula koizumii–N. kamtschatica* Zone are common to abundant and overall show moderate to good states of preservation (Table 2). Diatoms of the upper Pliocene to Pleistocene *N. koizumii* and *Actinocyclus oculatus* Zone show more

Lithologic unit	Diatom zone	Hole 794A Core, section, interval (cm)	Depth (mbsf)	Thalassionema nitzschiodies	Thalassiosira antiqua	Т. сопvеха	T. eccentrica	T. gravida	T. leptopus	T. marujamica	T. miocenica	T. nidutus	T. nordenskioldi	T. oestrupii	T. trifulta	T. zabelinae	Thalassiothrix longissima	Sponge spicules	
	N. koizumii	8H-1, 124–125 8H-2, 124–125 8H-3, 124–125 8H-4, 124–125 8H-6, 124–125	65.04 66.54 68.04 69.54 72.54	C R F					F					R		e	R	R R F F	
IB	N. koi–N. kamt.	9H-1, 124-125 9H-2, 124-125 9H-3, 124-125 9H-4, 124-125 9H-5, 124-125 9H-0, 124-125	74.54 76.04 77.54 79.04 80.54 82.04	FCFRC		-		F R R	R R R			R		R R F R		R R R R	FFRF	RFFFF	O D T
		10H-1, 124-125 10H-2, 124-125 10H-3, 124-125 10H-4, 124-125 10H-5, 124-125 10H-6, 124-125	85.54 87.04 88.54 90.04 91.54	RFFC	D			R	R R F			R	F	R R F	R R	RFR	R F F C R F	F R R F R F R F	z
IIA	Thaslassiosira oestrupii	11H-1, 124-125 11H-2, 124-125 11H-5, 124-125 11H-6, 124-125 12H-1, 123-125 12H-2, 123-125	95.54 95.04 99.54 101 103 104.5	FFCF	R R			r R R					r R	R R F		F C R	F C F F F F F	FFFRF	
		12H-3, 124–125 12H-4, 107–108 12H-5, 124–126 12H-6, 124–126	106 107.4 109 110.5	F C C				R F					R	R			F F C	R R R	

variable states of abundance and preservation. Similar to the assemblages in Hole 794A, Hole 795A assemblages are dominated by subtropical to subarctic planktonic diatom assemblages, with rare to few occurrences of benthic or nearshore diatoms such as *Diploneis bombus*, *D. smithii*, and *Actinoptychus senarius* (Table 2).

C. marginatus is much less common in Hole 795A above Sample 127-795A-16H-3, 120–121 cm, at 146.50 mbsf (Tables 2 and 4, Fig. 3). Although there are common occurrences of *C. marginatus* above 146.50 mbsf in Samples 127-795A-15H-5, 120–121 cm, and -15H-4, 50–52 cm, we generally recognize the base of the ODTZ between Samples 127-795A-16H-4, 74–79 cm (see radiolarian data), and 127-795A-16H-3, 120–121 cm, at 147.54–146.50 mbsf (Table 4), at the last consistent occurrence of common *C. marginatus* (Fig. 3). In addition to Sample 127-795A-16H-3, 120–121 cm, Samples 127-795A-16H-4, 115–116 cm (147.95 mbsf), and -18H-1, 120–121 cm (154.00 mbsf) also show common girdle bands of *C. marginatus* (Fig. 3). Above this horizon, silt-sized quartz grains are more evident.

In Sample 127-795A-13H-4, 120–121 cm, *C. marginatus* occurs with less frequency and the abundance of diatoms as a whole decreases (Table 2). Corresponding with this trend are higher abundances of sponge spicules (Zolnick et al., this volume; Pl. 2, Fig. 2).

Hole 797B

In Hole 797B, the base of the ODTZ as marked by the uppermost common occurrence of *C. marginatus*, occurs between Samples 127-797B-13H-3, 22–24 cm (113.62 mbsf), and -13H-2, 118–120 cm (113.08 mbsf) (Table 4, Figs. 3–4, Pls. 1–3). Below this depth diatoms from the *N. koizumii* and *N. koizumii–N. kamtschatica* Zone are mostly com-

mon to abundant but fluctuate with intervals containing fewer diatoms (Table 3). The assemblages below the base ODTZ are dominated by *C. marginatus, C. radiatus, N. koizumii, N. kamtschatica, Thalassionema nitzschioides, and Thalassiosira zabelinae* (Table 3).

The top of the ODTZ in Hole 797B is between 102.70 and 101.75 mbsf (Table 4), in the slides examined for diatoms. The top of the zone was recognized in Sample 127-797B-12H-1, 22–24 cm where *C. marginatus* is less abundant (Table 3 and Fig. 3). As in Hole 794A, *C. marginatus* girdles are abundant within the transition zone at Hole 797B, probably as a result of extreme dissolution and increased downslope transport. Table 3 and Figure 4 (see discussion in the radiolarian section) also indicate the presence of common siliceous sponge spicules near the top of the ODTZ in Samples 127-797B-12H-2, 22–24 cm (102.62 mbsf), and -12H-3, 22–24 cm (104.12 mbsf).

Radiolarians

Semiquantitative estimates of the abundance of five different types of siliceous microfossils which are >63 μ m in size (Fig. 4) show that the percent contribution of radiolarians (both whole and broken) is highly variable. In general, radiolarians increase between the base and the lower part of the opal dissolution transition zone. The upper portion of the ODTZ, particularly in Hole 795A, does not appear to have an increasing trend, and radiolarians are present in percentages (~40%) which are lower than those observed in samples from Holes 794A and 797B. However, sponge spicules are present in higher abundances in samples from Hole 795A, and act to dilute the percentage of radiolarians in Hole 795A. The increased numbers of siliceous

Neodenticula kamtschatica Coscinodiscus marginatus Hemidiscus cuneiformis Coscinodiscus radiatus Actinoptychus senarius C. marginatus (girdles) Stephanopyxis turris Rhizosolenia barboi Chaetoceros spore Azpeitia nodulifer Diploneis bombus Actinocyclus sp. Lithologic unit R. curvirostris Diatom zone Nitzschia sp. Preservation N. koizumii Abundance seminae Rouxia sp. D. smithii Hole 795A Core, section. Depth interval (cm) (mbsf) ž 12H-1, 120-121 105.50 С М R R P R 12H-2, 120-121 107.00 R 12H-3, 120-121 108.50 R P P 12H-5, 120-121 111.50 R R 12H-6, 120-121 113.00 Μ F R R R F 13H-1, 120-121 115.00 R P R IB 13H-2, 120-121 116.50 Ē P F R R 13H-3, 120-121 118.00 R P R C F F 13H-4, 120-121 119.50 C M F R 13H-5, 120-121 C M R R 121.00 R F F F F G R R R 13H-6, 120-121 122.50 A F R R C 13H-7, 61-63 123.41 M R M F F R R R 14H-2, 120-121 126.00 C R R N. koizumii R 14H-3, 120-121 P R R 127.50 F 129.00 С M R R R R 14H-4, 120-121 F R F 14H-5, 120-121 130.50 C G FR F 14H-6, 120-121 132.00 C G R R R R 14H-7, 62-63 132.92 M 15H-1, 120-121 134.00 С 15H-2, 120-121 135.00 M R 15H-4, 120-121 140.00 G С R R R F F 15H-5, 120-121 141.50 M C C С A 16H-1, 120-121 143.50 Μ F A R F 16H-2, 120-121 145.00 C Μ F 146.50 16H-3, 120-121 C M F F 16H-4, 115-116 M C С 147.95 A 18H-1, 120-121 С С C F 154.00 M F F F F F F 18H-2, 120-121 155.60 C M 18H-3, 120-121 157.10 С M F F F 18H-5, 120-121 160.10 R P R 11 F R 18H-6, 120-121 161.60 С M R C 19H-1, 120-121 163.60 M C F A A C C 19H-2, 120-121 165.10 М F A R A R P 19H-3, 120-121 166.60 C C F F 19H-4, 120-121 168.10 M R R CCC M F 19H-5, 120-121 169.60 C M CRCCFCF R 19H-6, 120-121 C 171.10 N. koizumii -RCCFRCC C G R R 21X-1, 120-121 182.80 N. kamtschatica F 21X-2, 120-121 184 30 M M 21X-3, 120-121 185.80 A 21X-5, 120-121 188.80 A G 21X-6, 120-121 190.30 C Μ F 22X-1, 120-121 192.40 Μ F F A 22X-2, 120-121 193.90 М F R A C C 22X-3, 120-121 195.40 Μ F R A 22X-4, 120-121 196.90 C М F F 22X-5, 120-121 198.40 C Μ C R 22X-6, 120-121 199.90 A G F

Table 2. Abundance and preservation of Pliocene and Pleistocene diagnostic diatoms from Leg 127, Hole 795A, Japan Sea. Also shown is the abundance of sponge spicules throughout the core. Biostratigraphic zonation of the core is after Shipboard Scientific Party (1990c).

Abundances are abbreviated as follows: A = abundant, C = common, F = few, R = rare, and B = barren. For preservation, G = good, M = moderate, and P = poor.

sponge spicules in Hole 795A may be due to increased downslope transport at this site which would also explain the relatively high amount of microfossil reworking at this site (Shipboard Scientific Party, 1990c).

Percent whole *Coscinodiscus marginatus* generally decrease upsection within the opal dissolution transition zone, and is also found in higher abundances in Hole 795A. Percentages of broken *C. marginatus* also appear to decrease with increasing depth in the opal dissolution transition zone. However, there is a marked peak in the upper portion of this transition zone in Hole 797B. Trends are not obvious in the percent *C. marginatus* girdle records, but because the girdles appear to be heavily silicified, their presence seems to be an indicator of increased levels of dissolution.

Although these measurements (Fig. 4) represent only semiquantitative estimates of the relative contribution of the five main categories of siliceous microfossils, the general trends of decreasing percent *C. marginatus*, and fluctuating yet increasing percent radiolarians and siliceous sponge spicules upsection within the opal dissolution transition zone is observed. Since radiolarians and sponges are less susceptible to dissolution than are diatoms (Johnson, 1974), these data

Lithologic unit	Diatom zone	Hole 795A Core, section, interval (cm)	Depth (mbsf)	Thalassionema nitzschioides	Thalassiosira antiqua	Т. сопчеха	T. eccentrica	T. gravida	T. leptopus	T. marujamica	T. miocenica	T. nidulus	T. nordenskioldi	T. oestrupii	T. trifulta	T. zabelinae	Thalassiothrix longissima	Sponge spicules	
IB		12H-1, 120-121 12H-2, 120-121 12H-3, 120-121 12H-5, 120-121 12H-6, 120-121 13H-1, 120-121 13H-2, 120-121 13H-3, 120-121 13H-4, 120-121 13H-5, 120-121 13H-6, 120-121 13H-6, 120-121	105.50 107.00 108.50 111.50 113.00 115.00 116.50 118.00 119.50 121.00 122.50 123.41	RCP	R		R	F	F					R F			R	R R R F R F R F R F R F R F R F R F R F	
	N. koizumii	13H-7, 61-05 14H-2, 120-121 14H-3, 120-121 14H-4, 120-121 14H-5, 120-121 14H-6, 120-121 14H-7, 62-63 15H-1, 120-121 15H-2, 120-121 15H-4, 120-121 15H-4, 120-121 16H-1, 120-121 16H-2, 120-121	123.41 126.00 127.50 129.00 130.50 132.92 134.00 135.00 140.00 141.50 143.50 145.00 146.50	R C F F C C C C C C C	R F F R	R		R F R R	F F R					R R		R	R R R F F F F F F	R R R R R R C F F F C F F	C T Z
П	N. koizumii – N. kamtschatica	101-4, 112-116 18H-1, 120-121 18H-2, 120-121 18H-5, 120-121 18H-6, 120-121 19H-1, 120-121 19H-1, 120-121 19H-1, 120-121 19H-1, 120-121 19H-4, 120-121 19H-6, 120-121 19H-6, 120-121 19H-6, 120-121 19H-6, 120-121 21X-1, 120-121 21X-2, 120-121 21X-5, 120-121 22X-1, 120-121 22X-2, 120-121 22X-3, 120-121 22X-4, 120-121 22X-5, 120-121 22X-5, 120-121	14.95 154.00 155.60 157.10 160.10 161.60 163.60 165.10 166.60 168.10 169.60 171.10 182.80 184.30 184.30 185.80 184.30 192.40 193.90 195.40 196.90	F F F F F F F C C C F C C	R F			F	F					R R F		F F F F C C C C A C C	R R F F	FFFFRF FRFFFRRFFRRRRR FFFRRRRRR FFFRRRRRR	

suggest that the siliceous assemblages over this interval were subjected to differing degrees of dissolution which, in general, increased with decreasing age. The diatomaceous intervals below the base of the opal dissolution transition zone contain high percentages of all three forms of *Coscinodiscus marginatus*, which serve as a diluting component in the slides making the relative contribution of radiolarians to the biosiliceous assemblages quite low (Pl. 1).

ODTZ

The opal dissolution transition zone is located by combining the information obtained from observations of slides prepared for diatom and radiolarian paleontological analyses (Figs. 3–4 and Pls. 1–3). The

transition is very obvious in samples prepared for diatoms and radiolarians and occurs over the length of several cores. The ODTZ is thinnest (9.6 m) in Hole 794A, and thickest (28 m) in Hole 795A (Table 4), and the time represented by these thicknesses is ~400,000 yr. The dissolution transition zones appear to be synchronous between Sites 795 and 797 (2.7–2.3 Ma, Table 4) while it is found in slightly older sediments in Hole 794A (2.9–2.6 Ma, Table 4). We believe that the opal dissolution transition zone is likely synchronous between all sites, and that the apparent diachroneity is caused by uncertain age control (probably in Holes 795A and 797B).

The opal dissolution transition zone roughly correlates to the transition from Unit I silty clays, to Unit II clayey diatom oozes at all three sites (794, 795, and 797) as defined in Tamaki, Pisciotto, and

Table 3. Abundance and preservation of Pliocene and Pleistocene diagnostic diatoms from ODP Leg 127, Hole 797B, Japan Sea. Also shown is the abundance of sponge spicules throughout the core. Biostratigraphic zonation of the core is after Shipboard Scientific Party (1990d).

Lithologic unit	Diatom zone	Hole 797B Core, section, interval (cm)	Depth (mbsf)	Abundance	Preservation	Coscinodiscus marginatus	C. marginatus (girdles)	Actinocyclus sp.	Actinoptychus senarius	Azpeitia nodulifer	Chaetoceros spore	Coscinodiscus radiatus	Diploneis bombus	D. smithü	Hemidiscus cuneiformis	Neodenticula kamtschatica	N. koizumii	N. seminae	Nitzschia sp.	Rhizosolenia barboi	R. curvirostris	Rouxia sp.
IB	N. koizumii	11H-1, 22-24 11H-2, 22-24 11H-3, 22-24 11H-4, 22-24 11H-5, 22-24 11H-6, 22-24 11H-7, 22-24 12H-1, 22-24 12H-1, 22-24 12H-4, 22-24 12H-5, 22-24 12H-6, 22-24 12H-6, 22-24 12H-7, 22-24 13H-1, 22-24 13H-3, 22-24 13H-3, 22-24 13H-5, 22-24 13H-6, 22-24 13H-6, 22-24 13H-7, 22-24 13H-7, 22-24 13H-6, 22-24 13H-6, 22-24 13H-7, 22-24 13	91.62 93.12 94.62 96.12 97.62 99.12 100.62 101.12 102.62 104.12 105.62 107.12 108.62 110.12 110.62 112.12 113.62 115.12 116.62 118.12 119.62	R R R A C F R F A C C F A F F A C C A C C A	P P G M M P P M M M G P P M M M M M G G	R F F F F F F F F C C C C C C F	F A F R R R R C C F F R	R F R	R R R	R R R R R R R R R R	F R R R F R R F R R F R F R	FFF FFFF FFF FFF FFF FFF	F R R	R R	F R R F F	R	C F R F F F F R C F R C C A F C	R F F R R		R F R R R R C R R R R R F F R R R R R R		R R
п	N. koizumii – N. kamtschatica	$\begin{array}{c} 14\text{H-1}, 23-25\\ 14\text{H-2}, 23-25\\ 14\text{H-3}, 23-25\\ 14\text{H-4}, 23-25\\ 14\text{H-5}, 23-25\\ 15\text{H-1}, 12-14\\ 15\text{H-2}, 22-24\\ 15\text{H-3}, 22-24\\ 15\text{H-5}, 23-25\\ 15\text{H-6}, 23-25\\ 15\text{H-6}, 23-25\\ 15\text{H-6}, 22-24\\ 16\text{H-1}, 22-24\\ 16\text{H-1}, 22-24\\ 16\text{H-2}, 22-24\\ 16\text{H-2}, 22-24\\ 16\text{H-5}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-7}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-7}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-7}, 22-24\\ 16\text{H-7}, 22-24\\ 16\text{H-7}, 22-24\\ 17\text{H-1}, 20-22\\ 17\text{H-2}, 3-5\\ 17\text{H-3}, 6-8\\ 17\text{H-4}, 22-24\\ 17\text{H-5}, 23-25\\ 17\text{H-6}, 23-25\\ 17\text{H-7}, 22-24\\ 18\text{H-1}, 22-24\\ 18\text{H-2}, 22-24\\ 18\text{H-4}, 22-24\\ 18\text{H-4}, 22-24\\ 18\text{H-4}, 22-24\\ 18\text{H-6}, 22-24\\ 18\text$	$\begin{array}{c} 120.13\\ 121.63\\ 123.13\\ 124.63\\ 126.13\\ 129.52\\ 131.12\\ 132.62\\ 134.12\\ 135.63\\ 137.13\\ 138.62\\ 139.12\\ 140.62\\ 142.12\\ 143.62\\ 145.12\\ 146.62\\ 148.12\\ 148.6\\ 149.93\\ 151.46\\ 152.92\\ 154.63\\ 156.13\\ 157.62\\ 158.12\\ 159.62\\ 161.12\\ 162.62\\ 164.12\\ 165.62\\ \end{array}$	C C C C C C A A C C C C F F F F F F F C C F F F F	M M M M M M M M M M M M P P P P P M M M P P P P M	C C C C C C C F C C C F F F F F F F F C C F F F F F C C A A C F	F F R R C A A F F F F F F F F F F F C C F F F F		R F R R R		R R R F F	R R F R R R R R R R	R F F	R R R	F R F F R R F F	R FFCC FF R R FF R C	F F F			RR RR RR R R R R R R R R R R R R R R R		R

Abundances are abbreviated as follows: A = abundant, C = common, F = few, R = rare, and B = barren. For preservation, G = good, M = moderate, and P = poor.

Lithologic unit	Diatom zone	Hole 794A Core, section, interval (cm)	Depth (mbsf)	Stephanopyxis turris	Thalassionema nitzschioides	Thalassiosira antiqua	Т. сопvеха	T. eccentrica	T. gravida	T. leptopus	T, marujamica	T. miocenica	T. nidulus	T. nordenskioldi	T. oestrupii	T. trifulta	T. zabelinae	Thalassiothrix longissima	Sponge spicules	
IB	N. koizumii	11H-1, 22–24 11H-2, 22–24 11H-3, 22–24 11H-5, 22–24 11H-6, 22–24 11H-6, 22–24 12H-1, 22–24 12H-1, 22–24 12H-3, 22–24 12H-4, 22–24 12H-5, 22–24 12H-6, 22–24 12H-6, 22–24 13H-1, 22–24 13H-1, 22–24 13H-3, 22–24 13H-4, 22–24 13H-5, 22–24 13H-6, 22–24 13H-6, 22–24 13H-6, 22–24 13H-6, 22–24	91.62 93.12 94.62 96.12 97.62 99.12 100.62 101.12 102.62 104.12 105.62 107.12 108.62 110.12 110.62 112.12 113.62 115.12 115.62 115.12 116.62	R R F R R R R	R R R C F F F F F F R C F R F F C F C F	R F R F R	R R R R	R F R	R R R R R R R R R R R R R R R R	F F F R F					R R R R R R R R R		R F F	R R R R R R R R R R R R R R R R R R R	R R R F F F R F C C F F F F F F F R R R	C D T Z
п	N. koizumii – N. kamtschatica	$\begin{array}{c} 14\text{H-1}, 23-25\\ 14\text{H-2}, 23-25\\ 14\text{H-3}, 23-25\\ 14\text{H-4}, 23-25\\ 14\text{H-5}, 23-25\\ 15\text{H-1}, 12-14\\ 15\text{H-2}, 22-24\\ 15\text{H-3}, 22-24\\ 15\text{H-5}, 23-25\\ 15\text{H-6}, 23-25\\ 15\text{H-6}, 23-25\\ 15\text{H-6}, 23-25\\ 15\text{H-7}, 22-24\\ 16\text{H-2}, 22-24\\ 16\text{H-2}, 22-24\\ 16\text{H-3}, 22-24\\ 16\text{H-4}, 22-24\\ 16\text{H-5}, 22-24\\ 16\text{H-5}, 22-24\\ 16\text{H-5}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-7}, 22-24\\ 16\text{H-7}, 22-24\\ 16\text{H-6}, 22-24\\ 16\text{H-7}, 22-24\\ 17\text{H-5}, 23-25\\ 17\text{H-7}, 22-24\\ 18\text{H-1}, 22-24\\ 18\text{H-3}, 22-24\\ 18\text{H-4}, 22-24\\ 18\text{H-5}, 22-24\\ 18\text{H-6}, 22-24\\$	120.13 121.63 123.13 124.63 126.13 129.52 131.12 132.62 134.12 135.63 137.13 138.62 149.12 140.62 142.12 140.62 142.12 146.62 148.12 148.6 149.93 151.46 152.92 154.63 156.13 157.62 158.12 159.62 161.12 165.62	R R R R R R R R R R R R R R R R R R	FFFFFFCCRFFFRRRRRRFFR RRRFCCCFR		R R	R	F F F F F F R R R R R R R F F F	F F F F F R R					R R		C F F R R	R R R R R R R R R	R R F F F F F F F F F F F F F F F F F F	

Allan, et al. (1990). The best correspondence of the opal dissolution transition zone with the lithologic units described during Leg 127 occurs in Hole 795A. The top of the transition zone in Hole 795A is found in Core 127-795A-13H-4 (Table 4). This level corresponds almost exactly to the top of Unit II which is distinguished by transitional indistinct color banding (Shipboard Scientific Party, 1990c). Although the opal dissolution transition zone appears to straddle the boundaries between lithological Units I and II in Holes 794A and

797B, the zone correlates well to smear slide records. The opal dissolution transition zone can be used to precisely define the transition between these two units.

PALEOCEANOGRAPHIC INTERPRETATION

There are three possible places where opal dissolution could have occurred. In the water column, on the seafloor at the sediment/water

Table 4. Location by sample, depth, and age, of the top and bottom of the late Pliocene opal dissolution transition zone (ODTZ) in Sites 794, 795, and 797.

Top ODTZ	Bottom ODTZ	Thickness ODTZ (m)
Hole 794A		
-9H-3, 59–61 cm, to -9H-3, 124–125 cm 79.89–80.54 mbsf 2.6 Ma	-10H-4, 124–125 cm, to -10H-5, 69–71 cm 88.54–89.49 mbsf 2.9 Ma	9.6 m
Hole 795A		
-13H-4, 120–121 cm, to -13H-5, 74–76 cm 119.50–120.54 mbsf 2.3 Ma	-16H-3, 120–121 cm, to -16H-4, 74–76 cm 146.50–147.54 mbsf 2.7 Ma	28.04 m
Hole 797B		
-12H-1, 22–24 cm, to -12H-1, 117–119 cm 101.75–102.7 mbsf 2.3 Ma	-13H-2, 118–120 cm, to -13H-3, 22–24 cm 113.08–113.62 mbsf 2.6 Ma	11.87 m

Ages were calculated using the age models of Table 5 and Figure 2, and rounded to the nearest 100,000 yr.

interface, or within the sediments. Post-depositional diagenetic processes can be ruled out as the cause of siliceous microfossil dissolution and *Coscinodiscus marginatus* concentration in the late Pliocene and Pleistocene sediments of Japan Sea Sites 794, 795, and 797. As the opal A/CT transition migrates upward, deeper sections would start dissolving earlier, and if *C. marginatus* was concentrated by diagenetic dissolution, the percentages of *C. marginatus* would be higher deeper in the sections, and this is not the case.

Dissolution at the sediment/water interface cannot be ruled out entirely, because there is some evidence which suggests that dissolution on the seafloor was important during the deposition of the Pleistocene dark and light layers (Alexandrovich, this volume). However, since sediment accumulation (burial) rates of the dissolved (upper Pliocene and Pleistocene) assemblages are not significantly lower than the burial rates of the Miocene and lower Pliocene diatomaceous oozes, the substantial increases in dissolution of the assemblages probably did not occur at the sediment/water interface.

The increased dissolution could have occurred in the water column prior to deposition as a result of depletion of available dissolved silica and competition for silica by plankton. Because *Coscinodiscus marginatus* is a dissolution-resistant species (Burckle, Sturz, and Emanuele, this volume), and is no longer present in any significant abundance (<5% of siliceous material) in the upper Pliocene and Pleistocene samples, we must assume that productivity declined during the opal dissolution transition zone. This seems to imply that during intervals of diatomaceous ooze deposition, circulation of the Japan Sea was quite different than it is today.

The age of the opal dissolution transition zone corresponds to a time of global cooling and build up of Northern Hemisphere glaciers (Barron and Baldauf, 1989; Raymo et al., 1990). Although the timing of these two events might suggest that the depositional changes were caused by cooling, it seems unlikely that cooling could be the sole cause of the changes since diatoms can be very productive in cold waters. However, because low numbers of radiolarians/gm (and generally poorer preservation) found in the sediments of piston core RC12-379 (Morley et al., 1986) are correlated to glacial stages, cooling may be a partial cause of the decreased preservation, or apparent reduced productivity, of the Pleistocene Japan Sea.

Dissolution and decreased productivity of siliceous microplankton implies that the waters of the Japan Sea became depleted with respect to dissolved silica. Matoba (1984) showed the depth of the sill at the Tsugaru Strait decreased rather rapidly between the late Pliocene and Pleistocene. If the dissolved silica content of Pacific deep waters of the Pliocene were similar to today's, a reduction of the influx of Pacific water into the Japan Sea at the Tsugaru Strait could have caused concentrations of dissolved silica in the Japan Sea to decrease. In turn, decreased silica concentrations would serve to lower productivity and increase dissolution of siliceous microorganisms.

SUMMARY

Abundance and preservation of siliceous microfossils is strongly controlled by the environmental constraints at the time of deposition. One of the most critical controls is the concentration of silica in the bottom water that is, in turn, related to circulation of deep water and upwelling patterns. Two key driving forces in the movement of deep water are (1) cooling of water, necessary to drive thermohaline circulation; and (2) bathymetry, which determines the flow path of bottom water.

The record of the subtropical to subarctic diatom Coscinodiscus marginatus suggests it is an especially sensitive indicator of not only diatom productivity, but its physical robustness is useful in determining preservation of siliceous microfossil assemblages. Its peak abundance in upper Pliocene sediments of the Japan Sea is synchronous in Holes 795A and 797B and is estimated to be 2.4 Ma. This age agrees well with C. marginatus abundances in the North Pacific and its relationship to the migration of the subarctic front and oceanwide water mass restructuring events (Sancetta and Silvestri, 1986). This interpretation is further supported by diatom data from DSDP Sites 578-580 in the northwest Pacific by Koizumi (1986). The high abundance of C. marginatus marks the base of what we have termed the opal dissolution transition zone (ODTZ) and it appears to be a few 100,000 yr older in Hole 794A (2.9 Ma). We suggest that this diachronous distribution may be an artifact of dating methods and needs to be further investigated.

The peak abundance of *C. marginatus* between 2.9 and 2.5 Ma is followed by some distinctive changes in the opaline fossil assemblages. Generally speaking, in the 10–30 m overlying very abundant *C. marginatus*, diatom frustules become increasingly fragmented, resulting in numerous disarticulated girdle bands in the sediments. Radiolarians within this interval are generally more poorly preserved, and siliceous sponge spicules appear to show a slight increase in numbers. The base of the ODTZ roughly correlates to the cessation of diatom oozes and the increase of silty and clay-dominated sediments in the Japan Sea. This pattern of increasing dissolution after the peak of *C. marginatus* ceases to be a significant component of the siliceous microfossil assemblages.

Table 5. Age control points for Leg 127 Sites 794, 795, and 797.

Depth (mbsf)	Age (Ma)	Control point	Rate (m/m.y.)				
Site 794							
0.00.0.00							
0.00 0.00	0.72		22 600				
24.60	0.73	р	33.699				
31.10	0.90	р	38.325				
34.10	0.98	р	37.500				
54.60	1.66	p	30.147				
58.80	1.88	p	19.091				
75.40	2.47	p	28.136				
89.80	2.92	p	32.000				
92.30	2.99	n	35 714				
95 50	3.08	P	35 556				
08.80	2 19	P	22,000				
109.30	2.40	P	40.707				
108.20	3.40	р	42.727				
125.10	3.88	p	35.208				
126.70	3.97	P	17.778				
133.80	4.10	p	54.615				
135.90	4.24	p	15.000				
139.90	4.40	p	25.000				
140.40	4 47	P	7 143				
143.80	4 57	P	34 000				
149.50	4.77	P	22,500				
146.50	4.11	P	25.500				
216.90	6.40	d	41.963				
255.70	7.20	d	48.500				
293.50	7.90	d	54.000				
530.50	14.40	с	36.462				
540.00	16.00	f	5.937				
Site 795							
0.00 0.00							
35.00	0.73	p	47.945				
46.00	0.90	p	64,706				
48.60	0.98	n	32 500				
87.00	1.66	P	56 471				
07.00	1.00	P	46 364				
124.00	1.00	P	40.304				
134.00	2.50	d	58.8/1				
200.90	3.50	d	66.900				
327.20	5.30	d	70.167				
394.40	6.40	d	61.091				
481.10	9.00	d	33.346				
560.50	10.70	d	46.706				
605.70	11.60	C	50.222				
635.30	13.30	d	17.412				
Site 797							
0.00 0.00							
38.30	0.73	p	52.466				
78.60	1.66	p	43 333				
81.90	1.80	d	23 571				
84 60	1.99		32 750				
126.00	2.00	p	30,000				
120.00	2.92	р	39.808				
133.80	2.99	р	111.429				
143.60	3.08	р	108.889				
150.80	3.18	р	72.000				
167.80	3.40	p	77.273				
243.40	5.30	d	39.789				
350.00	7.20	d	56,105				
369.60	11.60	c	4 455				
439.10	14.40	0	24 821				
	1	8.7	24.0/				
407.50	17.10		20 140				

Paleomagnetic data are from Hamano et al. (this volume), and biostratigraphic data are from Shipboard Scientific Party (1990b, 1990c, and 1990e) with the exception of the LO of *Neodenticula kantschatica* (2.5 Ma control point) in Site 795 which is from Koizumi (this volume). P refers to paleomagnetic control point, and d, c, and f, to diatom datum, coccolith, and foraminifer level control point, respectively.



Figure 2. Age vs. depth of Sites 794, 795, and 797.

The complex tectonic setting of the Japan Sea, coupled with Neogene high latitude glaciations, resulted in a Pliocene through Pleistocene depositional setting that was particularly sensitive to water mass fluctuations. The decreasing sill depth of the Tsugaru Strait during the late Pliocene and Pleistocene, in response to lowering of sea level brought on by Northern Hemisphere glaciation, likely resulted in a reduction of the influx of Pacific water into the Japan Sea and, therefore, a reduction in dissolved silica contents in the Japan Sea. Thus, the dissolved siliceous microfossil assemblages observed at Sites 794, 795, and 797 record global temperature and circulation changes that were intensified in the marginal basin setting of the Japan Sea.

FLORAL LIST

Actinocyclus oculatus Jousé

Actinocyclus sp.

- Actinoptychus senarius (Ehrenberg) Ehrenberg
- Azpeitia nodulifer (Schmidt) Fryxell and Sims

Chaetoceros Ehrenberg

Coscinodiscus marginatus Ehrenberg

Coscinodiscus radiatus Ehrenberg

Diploneis bombus Ehrenberg

Diploneis smithii (Brebisson) Cleve

Hemidiscus cuneiformis Wallich

Neodenticula kamtschatica (Zabelina) Akiba and Yanagisawa

Neodenticula koizumii Akiba and Yanagisawa

Neodenticula seminae (Simonsen and Kanaya) Akiba and Yanagisawa



Figure 3. Variation in the relative abundance of *Coscinodiscus marginatus* and *C. marginatus* girdles in Pliocene–Pleistocene sediments from Hole 794A, 795A, and 797B. Note the peak in *C. marginatus* girdles at a depth slightly above the last major occurrence of *C. marginatus* frustules, in Holes 794A and 797B. Abundancea are abbreviated as follows: A = abundant, C = common, F = few, R = rare, B = barren.



Figure 4. Semiquantitative estimates of the breakdown of biosiliceous material on slides prepared for radiolarians in the Japan Sea opal dissolution transition zone. Radiolarian fragments were included in the %radiolarians category. Note differences in percent scales.

Nitzschia sp.

Rhizosolenia barboi (Brun) Tempere and Peragallo Rhizosolenia curvirostris Jousé Stephanopyxis turris (Grev. and Arn.) Ralfs Thalassionema nitzschioides (Grunow) Peragallo Thalassiosira antiqua (Grunow) Cleve-Euler Thalassiosira convexa Muchina Thalassiosira eccentrica (Ehrenberg) Cleve Thalassiosira gravida Cleve Thalassiosira leptopus (Grunow) Hasle and Fryxell Thalassiosira marujamica Sheshukova-Poretzkaya Thalassiosira nidulus (Tempère and Brun) Jousé Thalassiosira nordenskioldi Cleve Thalassiosira oestrupii (Ostenfeld) Proshkina-Labrenko Thalassiosira trifulta Fryxell Thalassiosira zabelinae Jousé Thalassiothrix longissima Cleve and Grunow

ACKNOWLEDGMENTS

Funding for this research was provided by JOI U.S. Science Support Program. This research would not have been possible without the laboratory assistance of Ann Melamed, Leslie Hewitt, and Laura Pound, all of San Francisco State University, and of Elaine Stock. Manuscript reviews by John Barron, Naja Mikkelsen, and Joe Morley are greatly appreciated.

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Date of initial receipt: 22 April 1991 Date of acceptance: 26 February 1992 Ms 127/128B-140



Photographs presented in the following plates were taken using a 35 mm camera mounted on a beam splitter on a Wild M20 microscope and a Zeiss Axioskop. Scale are indicated on the plates. EF refers to the England Finder location (Riedel and Foreman, 1961) of the specimen on the sample slide. Samples are kept at the Department of Geosciences, San Francisco State University, and the Department of Geology, Florida Atlantic University.

Plate 1. Photomicrographs from samples found below the opal dissolution transition zone. *Coscinodiscus marginatus* valves largely intact, showing little dissolution. Both whole girdles and valves largely intact. **1.** Sample 127-797B-13H-4, 22–24 cm. **2.** Sample 127-794A-11H-1, 60–62 cm, EF P19/3–4. **3.** Sample 127-797B-13H-4, 22–24 cm.





80 µm

40 µm



20 µm

Plate 2. Photomicrographs from samples found within the opal dissolution transition zone. Diatom valves show signs of dissolution and fragmentation (1 and 3) and samples prepared for radiolarians contain high abundances of siliceous sponge spicules (2). **1.** Sample 127-797B-12H-2, 22–24 cm. **2.** Sample 127-795A-13H-7, 51–53 cm, EF H10/4. **3.** Sample 127-794A-9H-6, 124–125 cm.



80 µm



80 µm



63

3

40 µm

Plate 3. Photomicrographs from samples found within the opal dissolution transition zone. Diatom valves are dissolved and fragmented (1, 2, and 3), and broken diatom girdles are abundant (1 and 2). 1. Sample 127-794A-10H-3, 124-125 cm. 2. Sample 127-797B-13H-2, 22-24 cm. 3. Sample 127-794A-9H-6, 59-61 cm, EF K12/3.