

*Benthic Mineral Cycling
in Marine Ecosystems:*

*Adriatic Sea
versus
North Sea*

Lutz Lohse



*Netherlands Institute for Sea Research
(NIOZ)*

Department of Marine Chemistry and Geology

Texel

The Netherlands

Lohse et al. Benthic Mineral Cycling, ITCP, Nov 1998

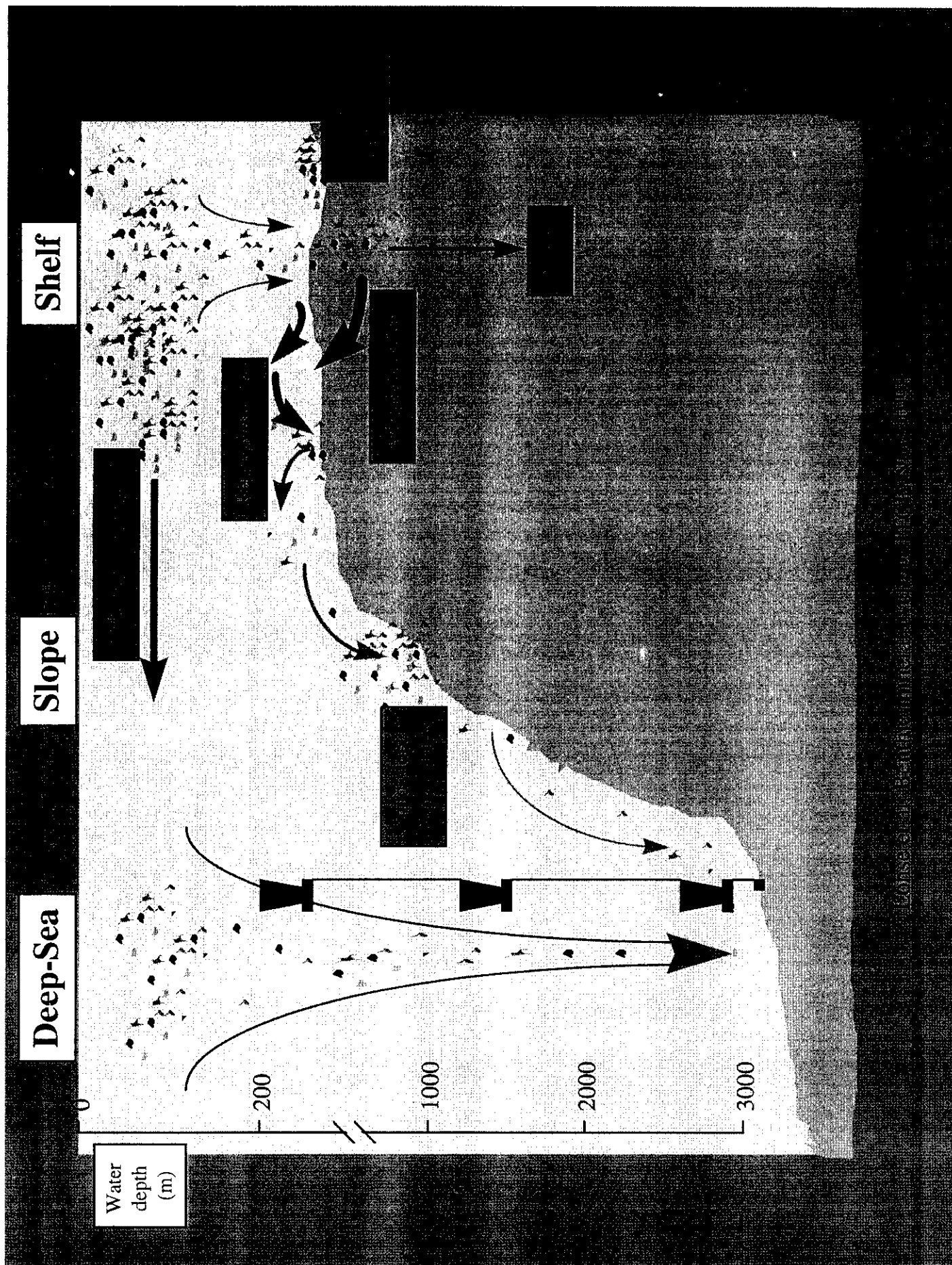
①

Outline

- *General introduction to benthic-pelagic coupling*
 - *sedimentation of organic matter in marine ecosystems*
 - *general benthic processes*
 - *early diagenesis*
 - *methodological considerations*

- *Case Studies*
 1. *Adriatic Sea*
 2. *North Sea*

 3. *N. W. European Continental Margin*



Importance of shelf sediments in global biochemical cycles

- Although the continental shelf covers only 15% of the world ocean it accounts for nearly 50% of the global primary production
- 85% of the global sediment mineralisation takes place on the shelf
- Benthic mineralisation accounts averagely for 45% (?) of the total mineralisation in shelf ecosystems

Typical values of transport and reaction parameters in marine sediments*

	Deep-Sea	Slope	Shelf/Coastal
water depth (m)	3000-6000	200-3000	<200
sedimentation rate (cm yr ⁻¹)	0.001	0.001-0.01	0-1
bioturbation (cm ² yr ⁻¹)	0.1	1.5	10
degradability (yr ⁻¹)	0.001	0.02	0.1-10
characteristic lenght scale (cm)	10-100	1-10	0.1-1
characteristic time scale (yr)	10000	100-1000	0.01-100

(*modified after Van Capellen and Gaillardet)

Methods to estimate (benthic) carbon mineralisation

1. Direct Methods

(Incubation of sediment or sediment-water enclosures)

Electron acceptor consumption or reduced product formation over time

- O_2 , N_2 , Fe_2^+ , CH_4^+ , ΣCO_2^-

Remineralisation of organic substances

- acetate, fatty acids, glucose

2. Indirect Methods

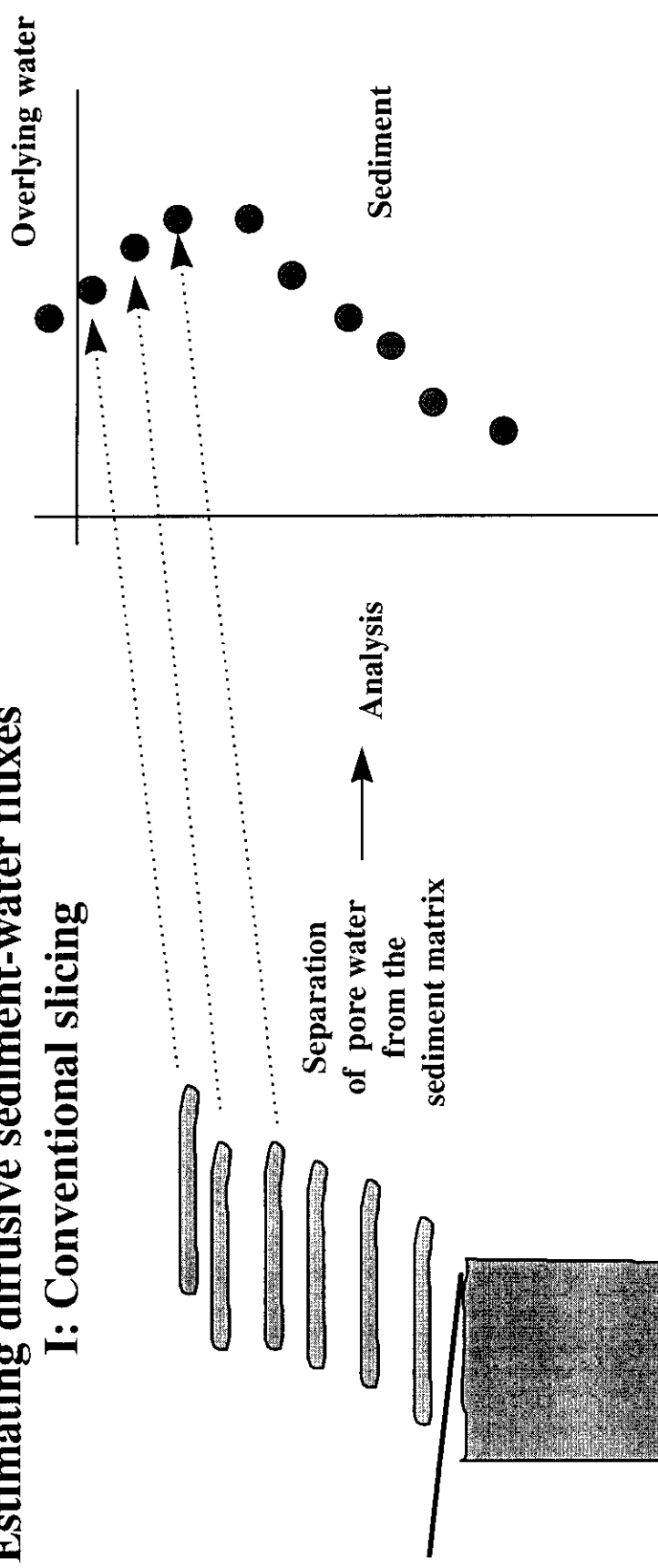
(Pore water profiling)

- solid phase organic carbon
- hydrolysable amino acids

- electron acceptors (e. g. O_2 , NO_3^-)
- reduced products (Mn_2^+ , Fe_2^+ , ΣCO_2^- , NH_4^+

Estimating diffusive sediment-water fluxes

I: Conventional slicing



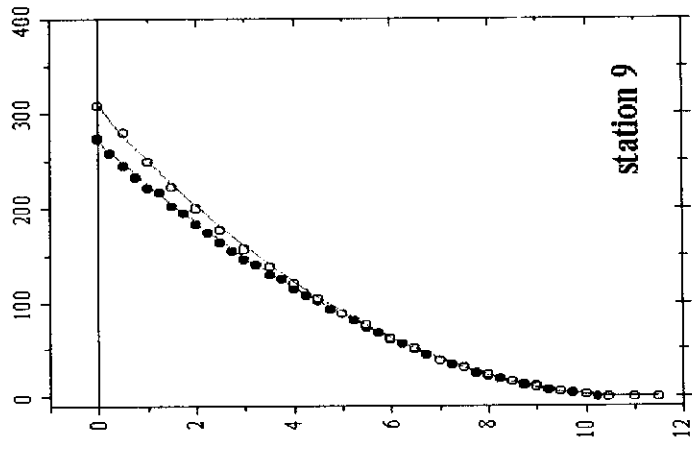
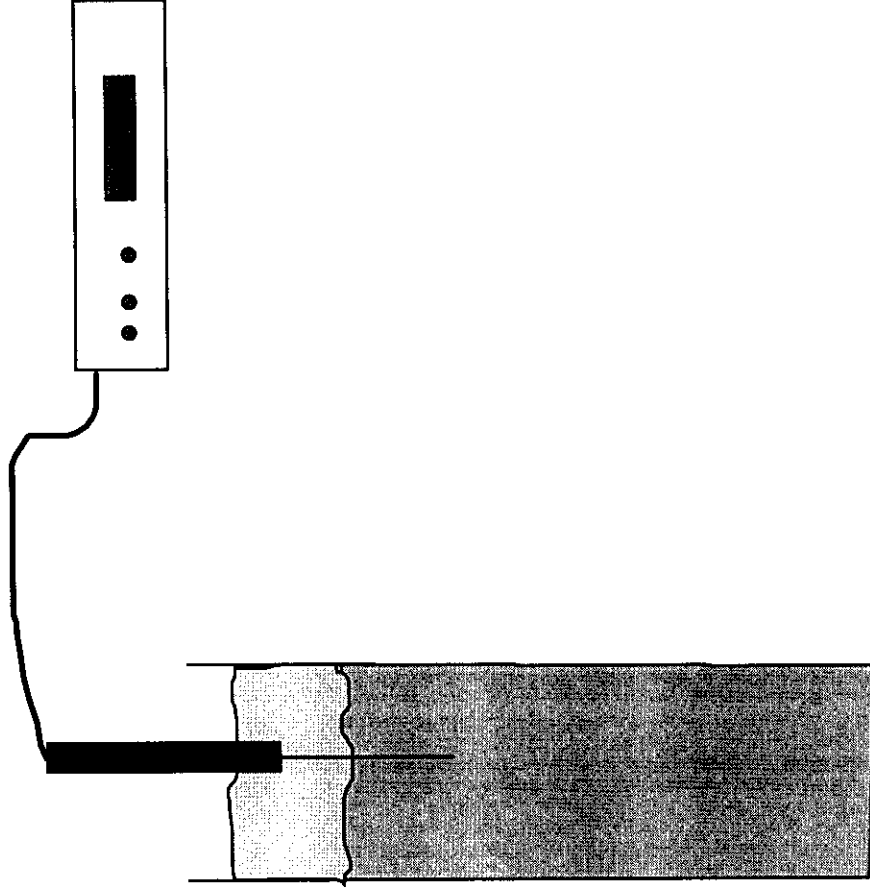
F (in $\mu\text{mol m}^{-2} \text{d}^{-1}$)

$$= D_s \times Dc/dz$$

+

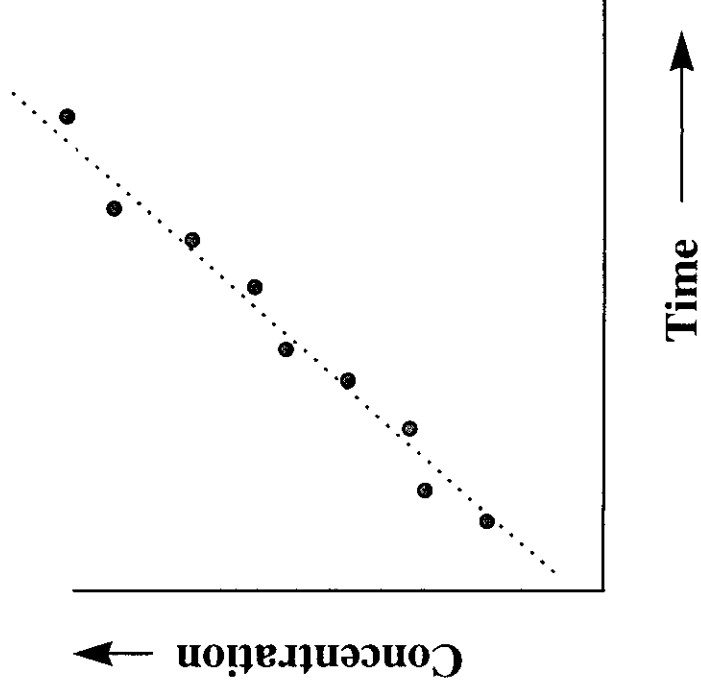
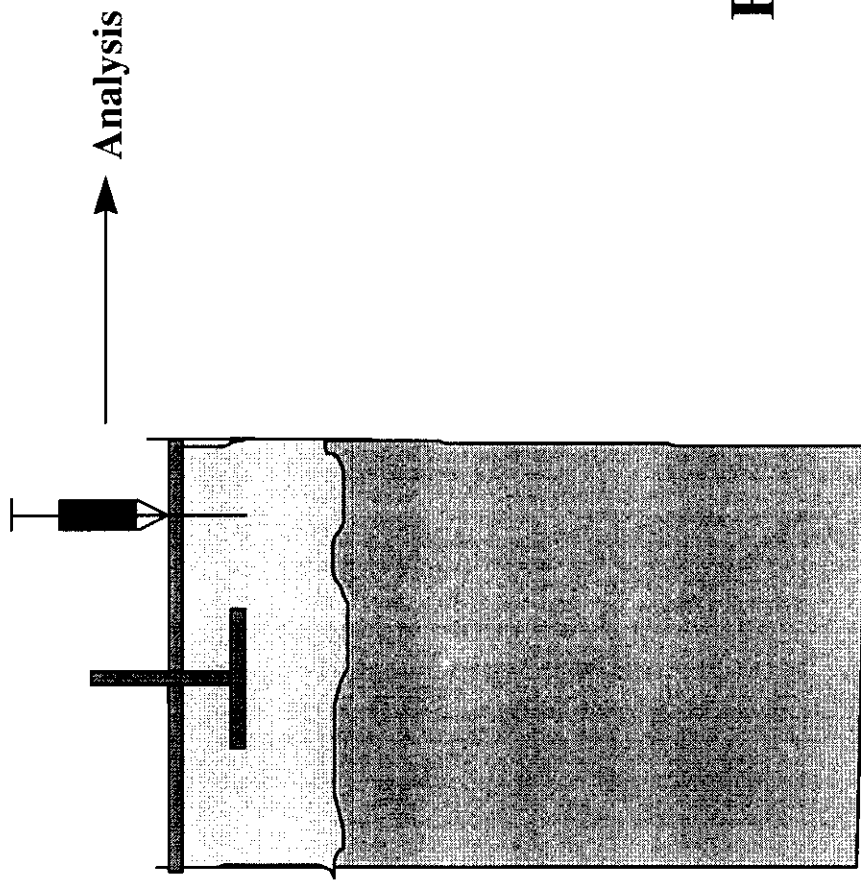
Estimating diffusive sediment-water fluxes

II: Ion-specific micro-electrodes

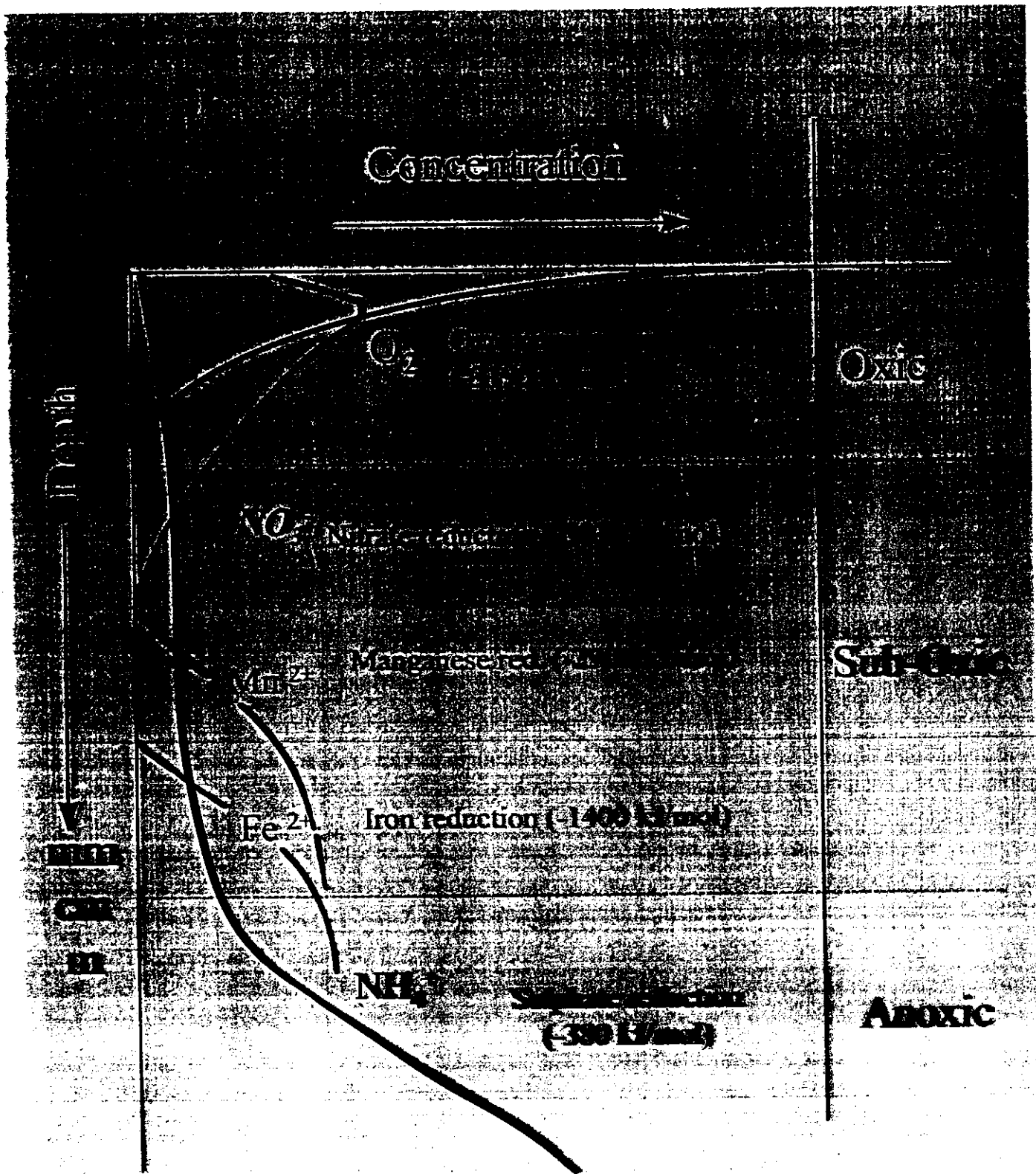


Estimating sediment-water fluxes

III. Whole core incubations

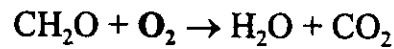


$$F = dc/dt \times A/V$$

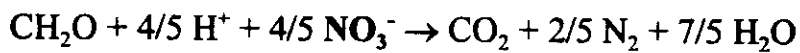


Organic carbon oxidation reactions and the subsequent oxidation of reduced species.

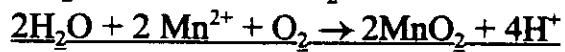
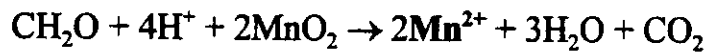
O₂ respiration:



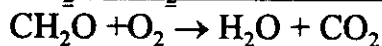
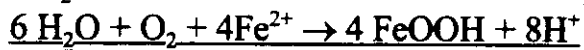
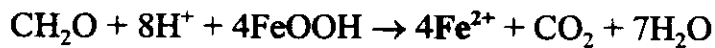
Denitrification:



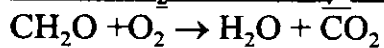
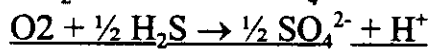
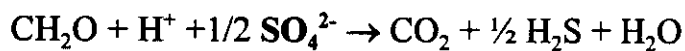
Mn reduction:



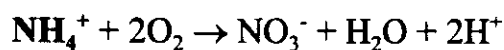
Fe reduction:



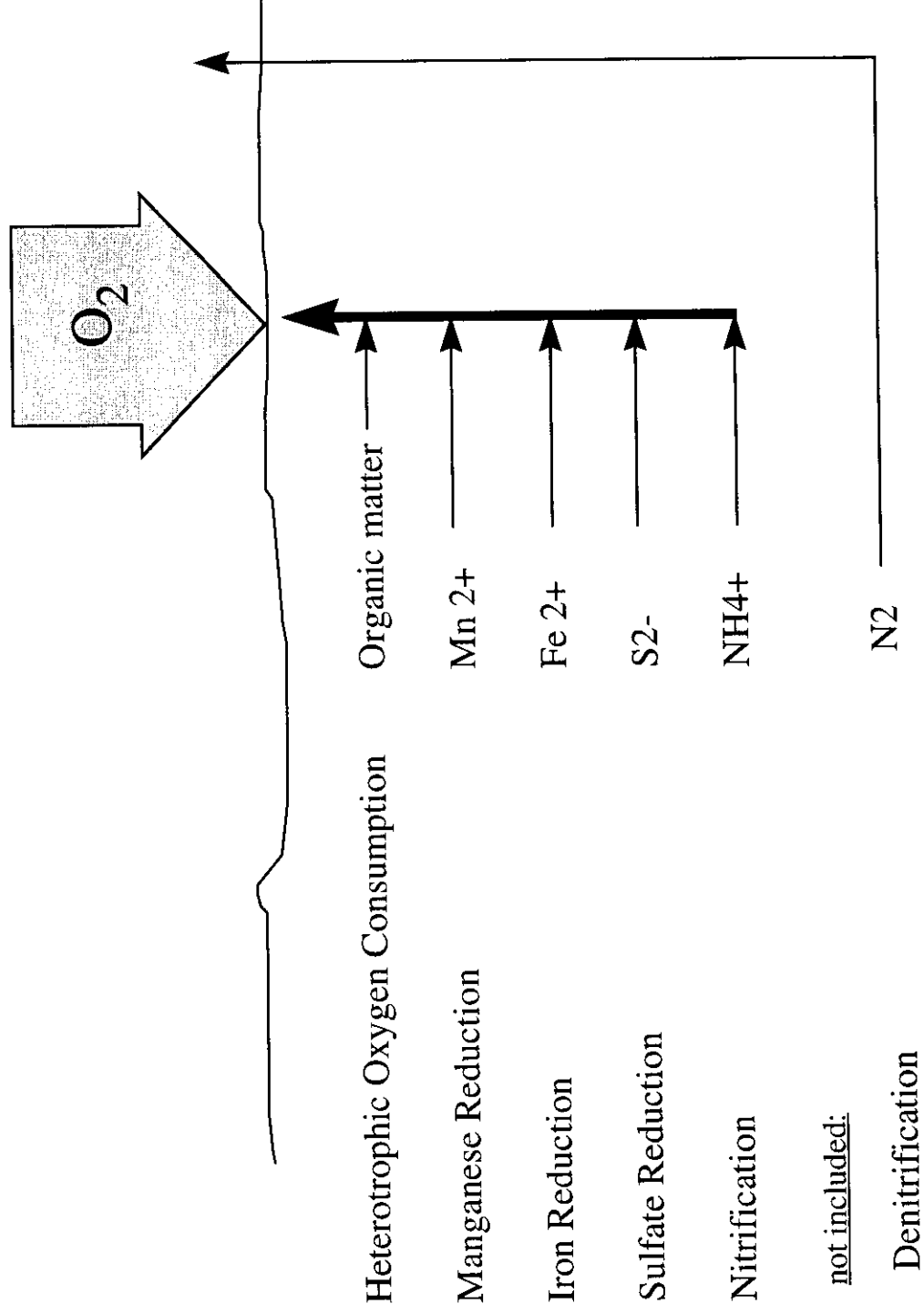
Sulfate reduction:



NH₄⁺ oxidation:



Carbon oxidation rates derived from benthic oxygen uptake rates



CARBON OXIDATION RATE

=

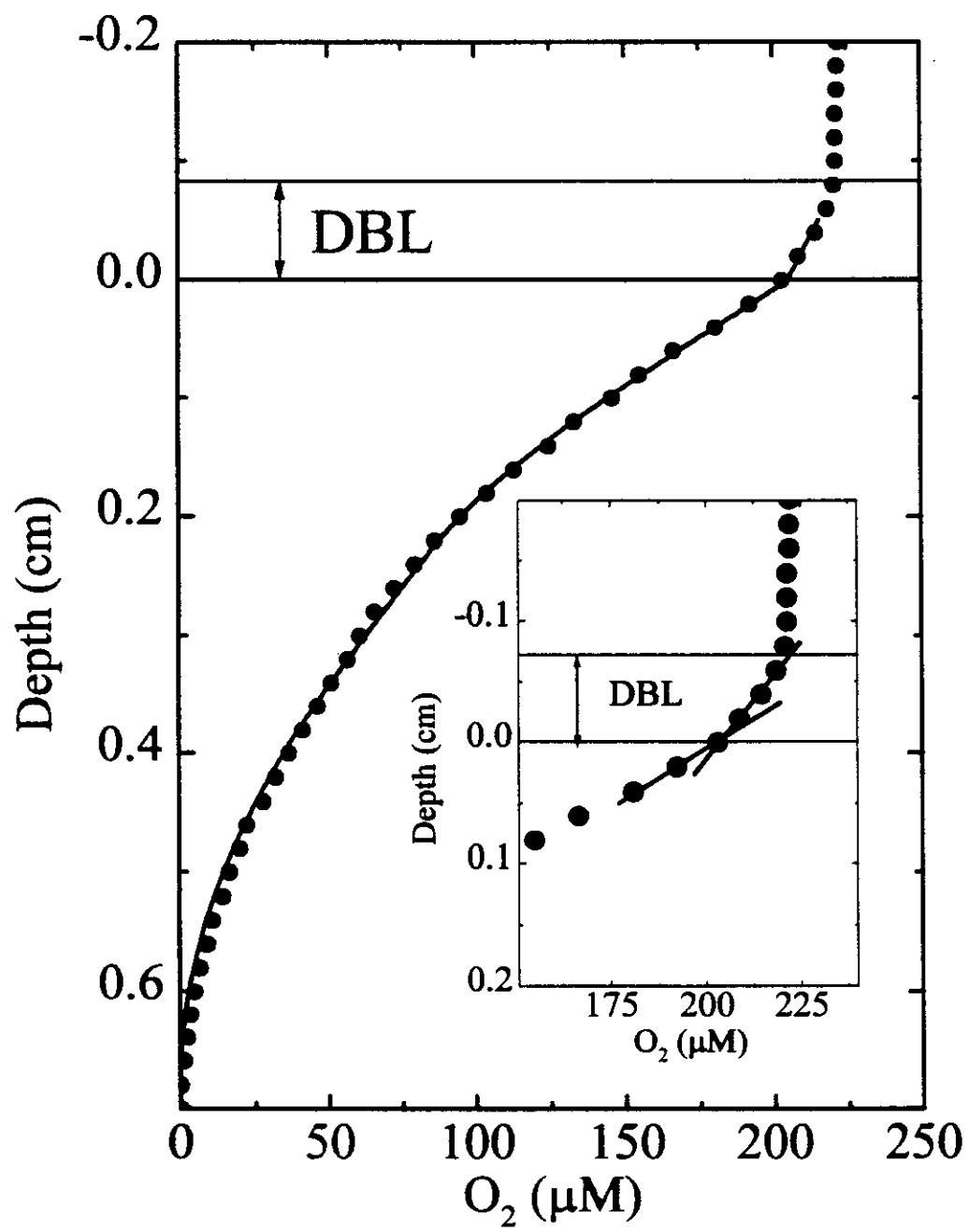
OXYGEN UPTAKE RATE

+ DENITRIFICATION

- OXYGEN CONSUMED BY NITRIFICATION

Constraints:

- 1) All reduced metabolites produced during suboxic and anoxic diagenesis are re-oxidised
- 2) CO₂ is the ultimate reaction product of all mineralisation processes
- 3) Burial of reduced compounds is negligible



Steady state distribution of dissolved solutes (oxygen)

$$0 = D_s \frac{d^2C}{dz^2} - R$$

Diffusion coefficient

Diffusive gradient

Reaction

15

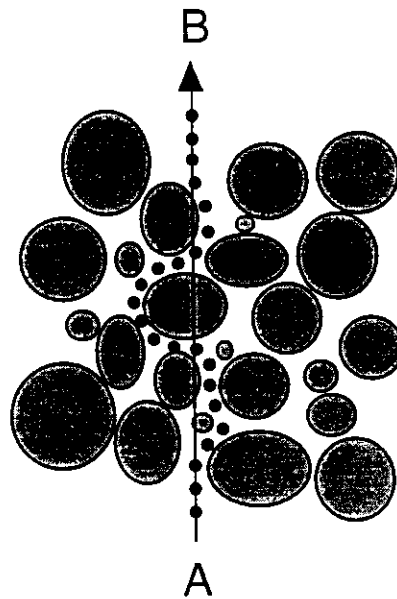


Figure 3.13. A solute trying to diffuse from point A to point B cannot pass through solid sediment particles (filled shapes). The solute must travel around these particles and so traverses the *tortuous* dotted path, rather than the direct solid-line path.

from Boudreau, 1998

Determination of the sediment diffusion coefficient D_s

$$D_s = \frac{D_0}{\phi F}$$

where D_0 is the diffusion coefficient for a free solution
 ϕ is the porosity
 F is the resistivity formation factor

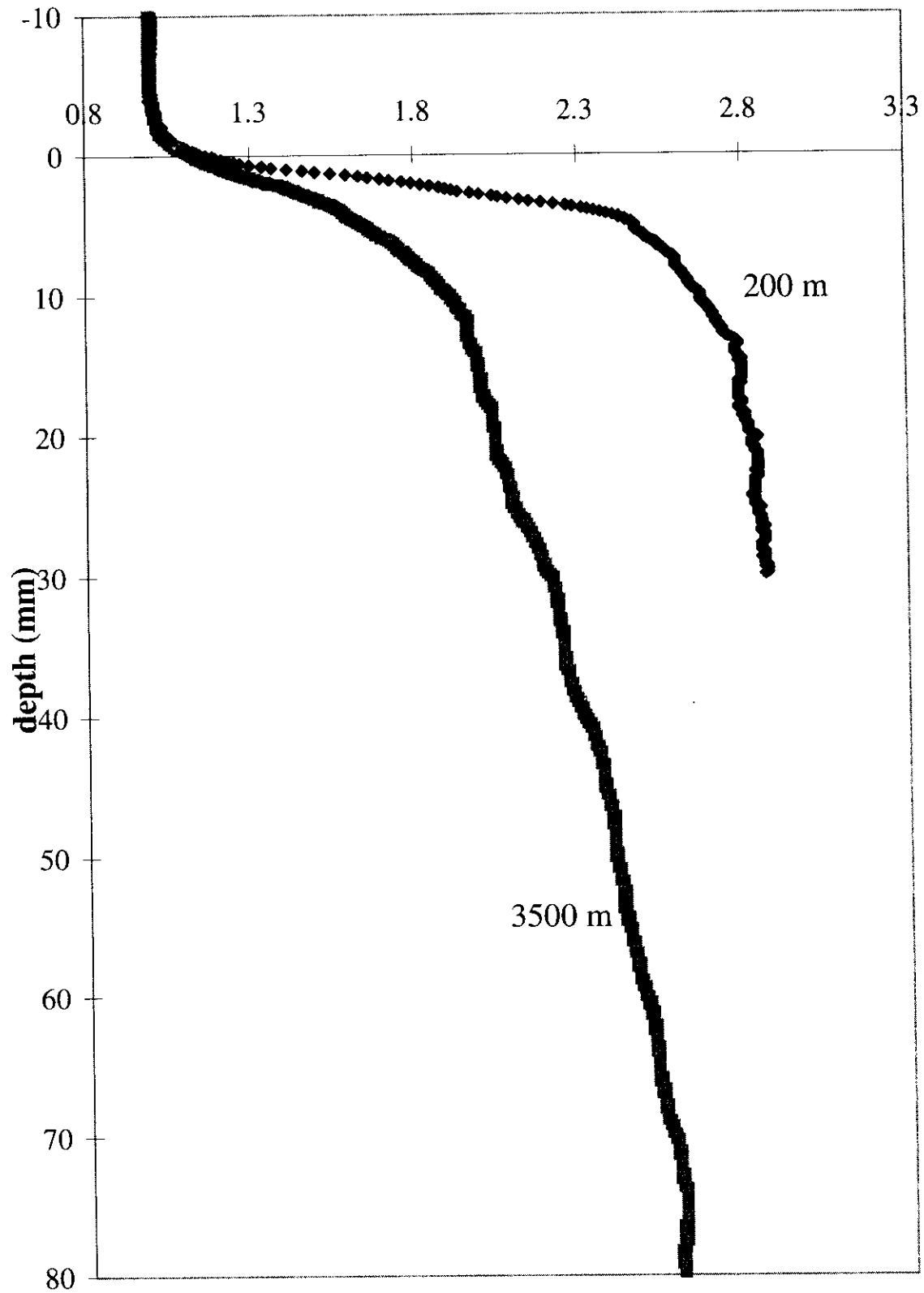
$$F = \frac{R_{\text{sediment}}}{R_{\text{water}}}$$

then the porosity profile can be calculated by

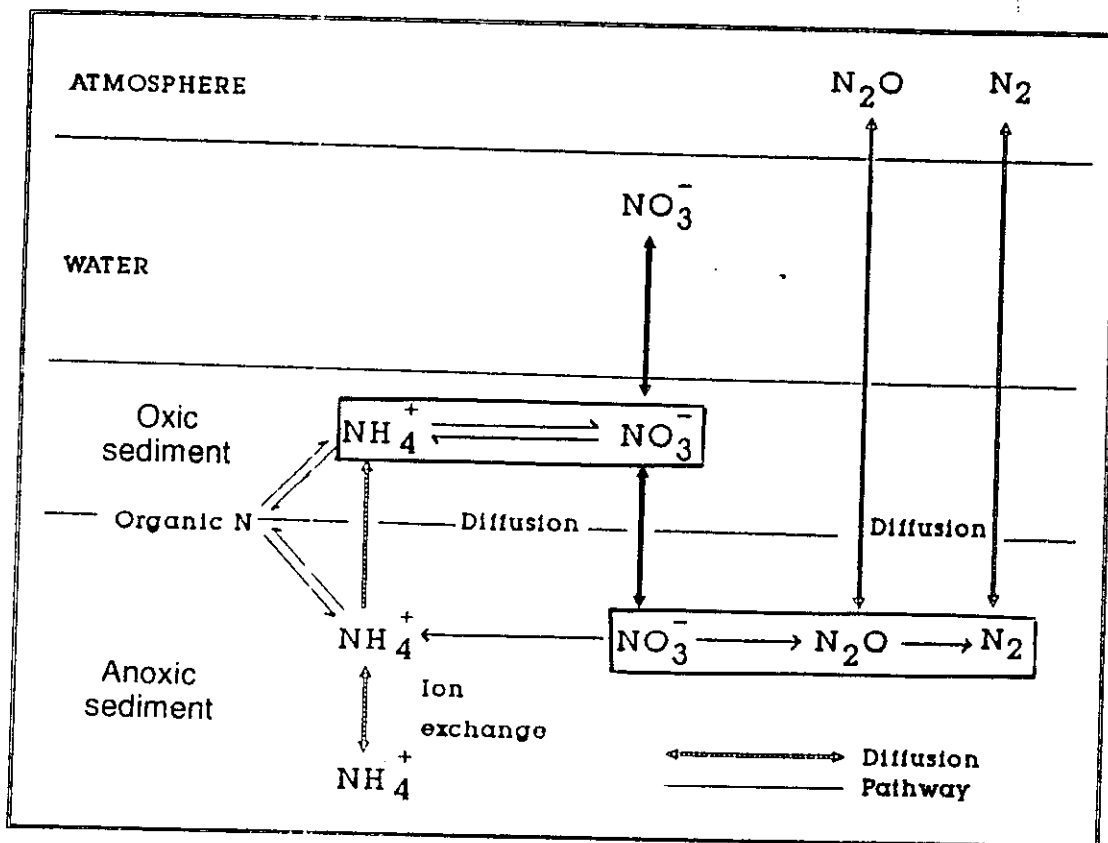
$$\phi = F^{\frac{1}{n}}$$

where n ranges between 1.5 and 3.5 for sandy and muddy sediments, respectively.

Resistivity Formation Factor F [-]

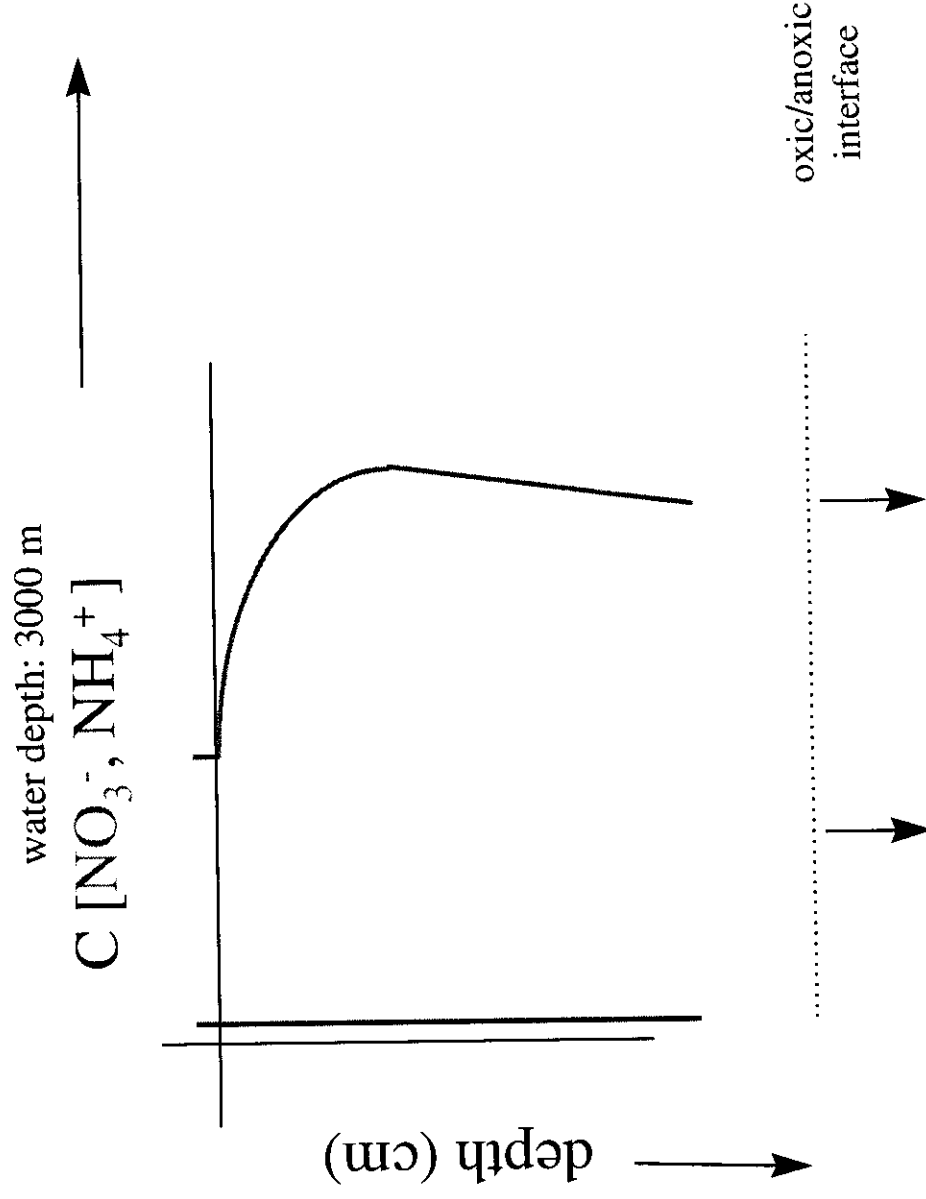


N-mineralisation in marine sediments: A conceptual view



Generalised nitrogen mineralisation

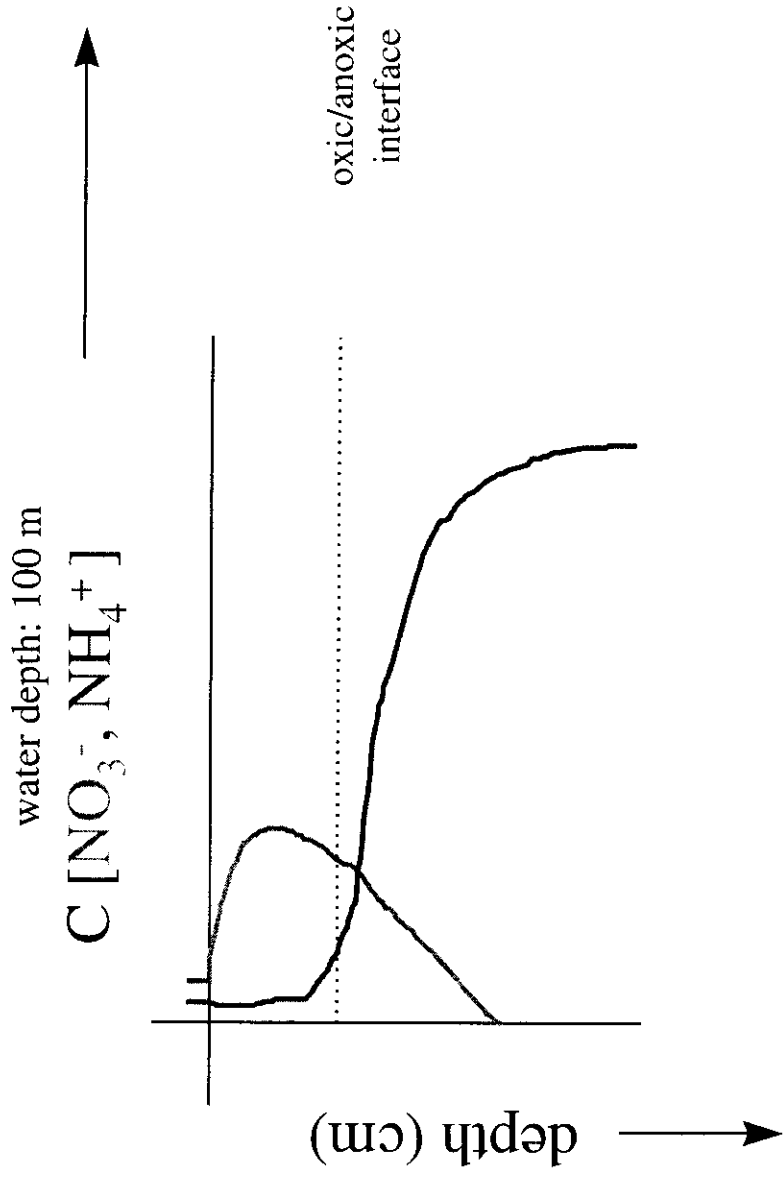
Case 1: Pelagic sediments



Lohse et al, Benthic mineral cycling, ITCP, Nov 1998

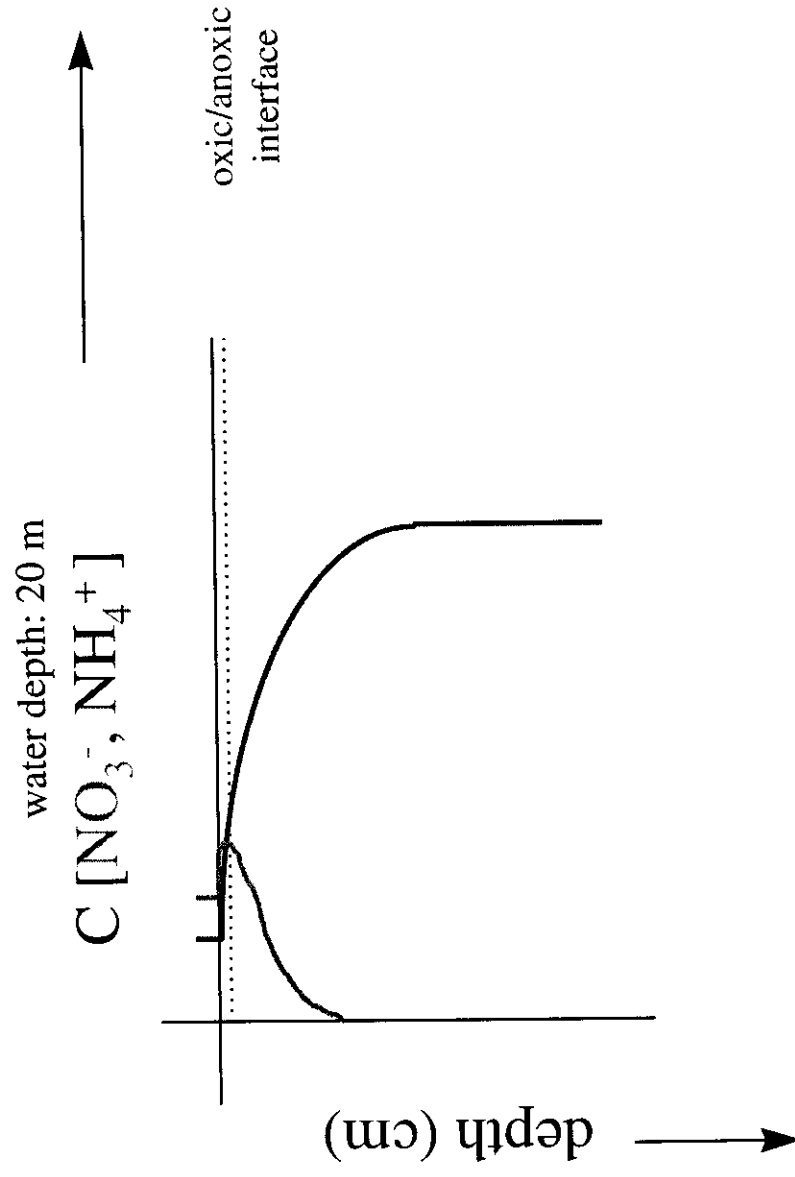
Generalised nitrogen mineralisation

Case 2: Shelf sediments



Generalised nitrogen mineralisation

Case 3: Coastal sediments



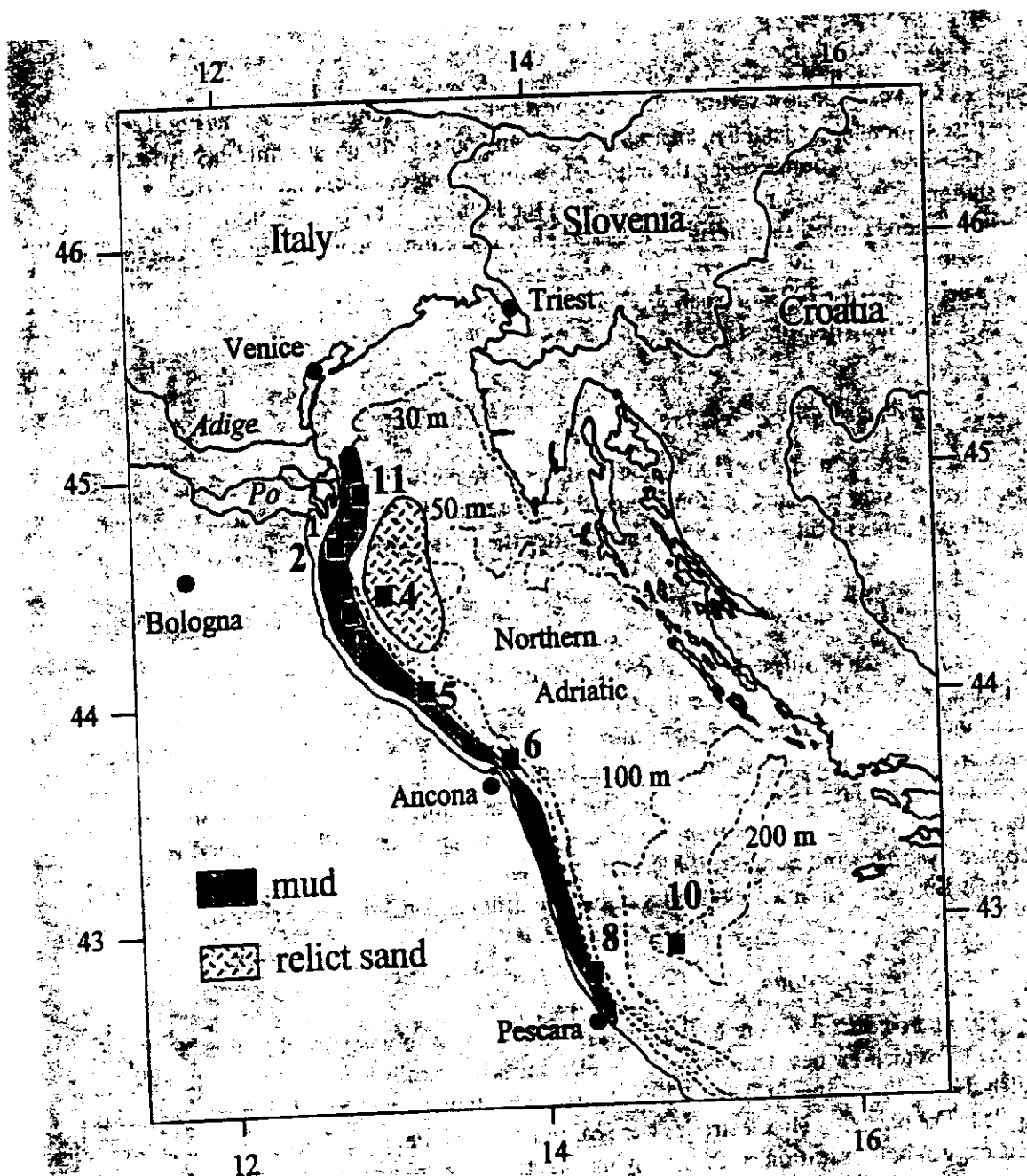
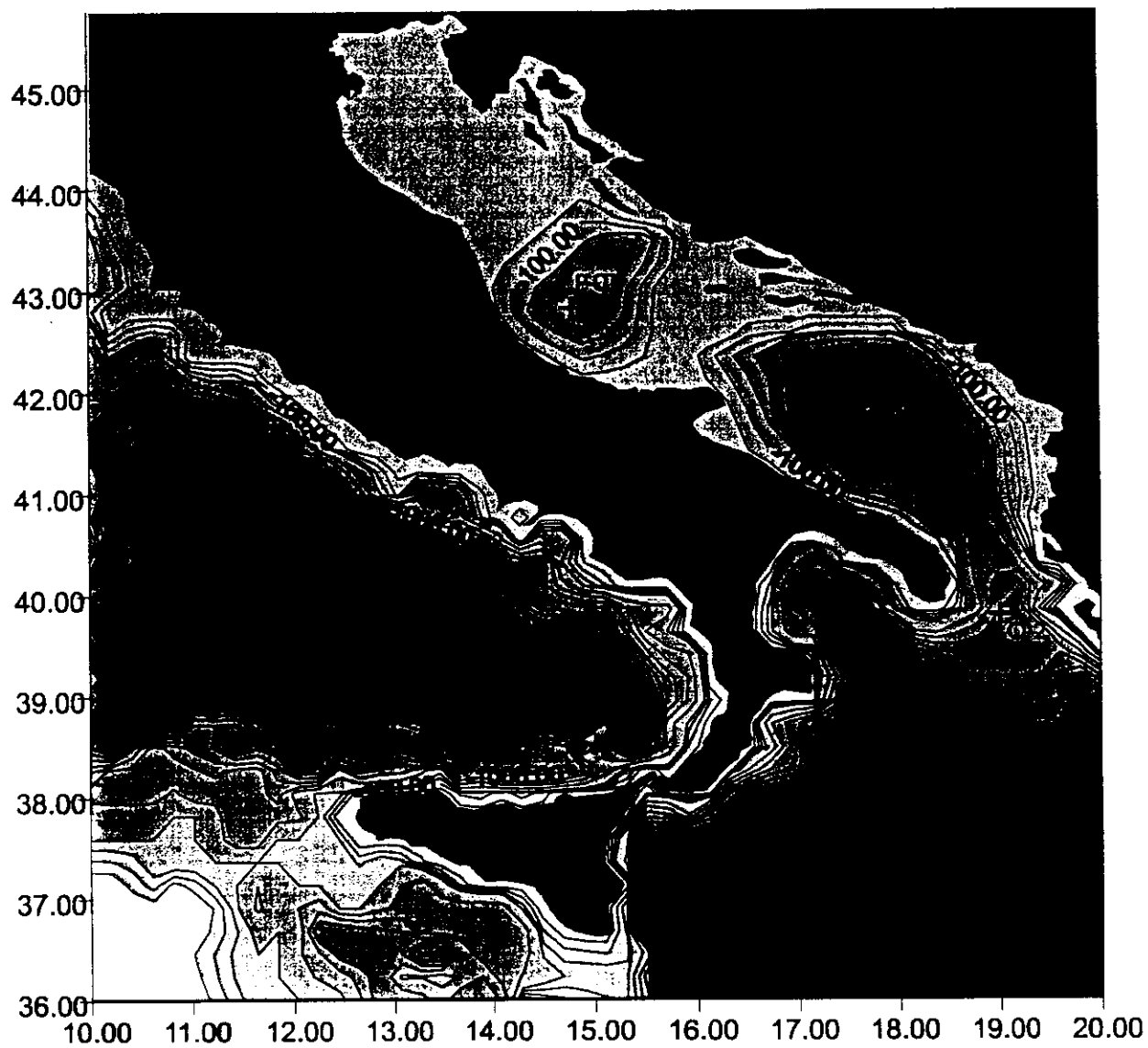


Fig. 1. Map of the Northern Adriatic Sea showing the stations occupied in March 1992 and August 1993.



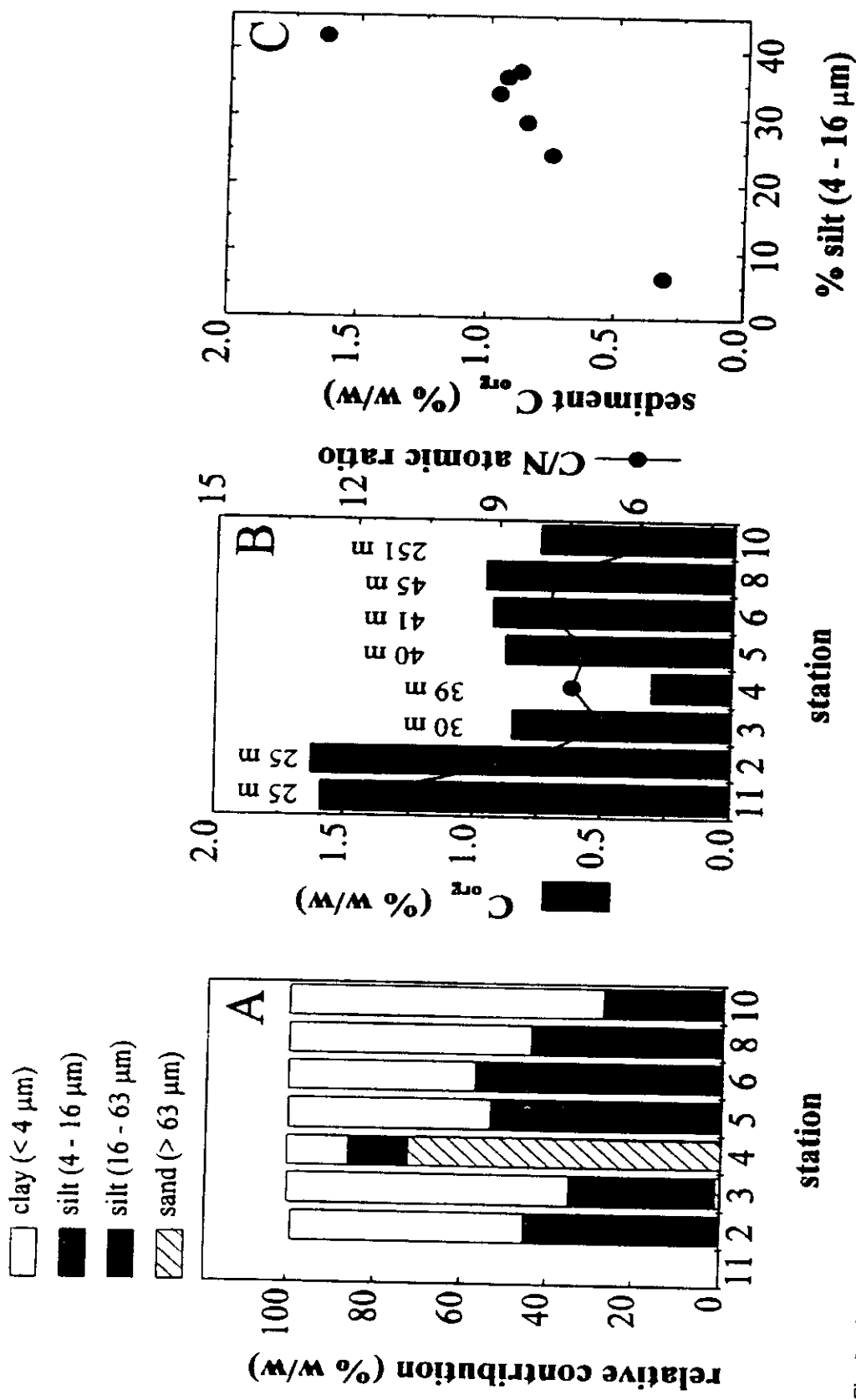


Fig. 2. Sediment characteristics of the occupied stations. (A) Sediment composition. Station 4 was located outside the coastal mudbelt in the relic sand area. Station 10 was located in the south at the Meso Adriatic Depression. Data were kindly provided by Dr P. Giordani, Institute for Marine Geology, Bologna. (B) Organic carbon content and the atomic C:N ratio for the surface 0-5 mm of the sediment. (C) Organic carbon content plotted against the silt (4-16 μm fraction) content of the sediments in August 1993.

Station	S (‰)		T (°C)		O ₂ (μmol dm ⁻³)		I _{sed} (μE m ⁻² s ⁻¹)	
	3-'92	8-'93	3-'92	8-'93	3-'92	8-'93	3-'92	8-'93
11		38.2		15.7		215 (88%)		15.2
2		38.1		14.1		192 (75%)		20.8
3	38.1	38.1	9.8	16.3	238 (85%)	207 (86%)	1.4	35.1
4		38.2		14.0		196 (77%)		8.1
5	38.2	38.2	10.0	14.0	277 (100%)	272 (108%)	-	19.4
6	38.3	38.3	10.5	13.1	275 (100%)	277 (108%)	1.1	11.4
8	38.3	38.1	11.2	15.7	255 (95%)	245 (102%)	0.5	-
10		38.3		10.9		224 (81%)		

Table 2. Bottom water salinity (S), temperature (T), oxygen concentration and PAR quantic irradiance at the seafloor, in March 1992 and August 1993. PAR values were kindly provided by prof. M. Innamorati and dr. L. Massi, Laboratorio di Ecologia, Florence, Italy.

Epping and Helder, 1998
Cont. Shelf Res 17 (4) 1737-1769

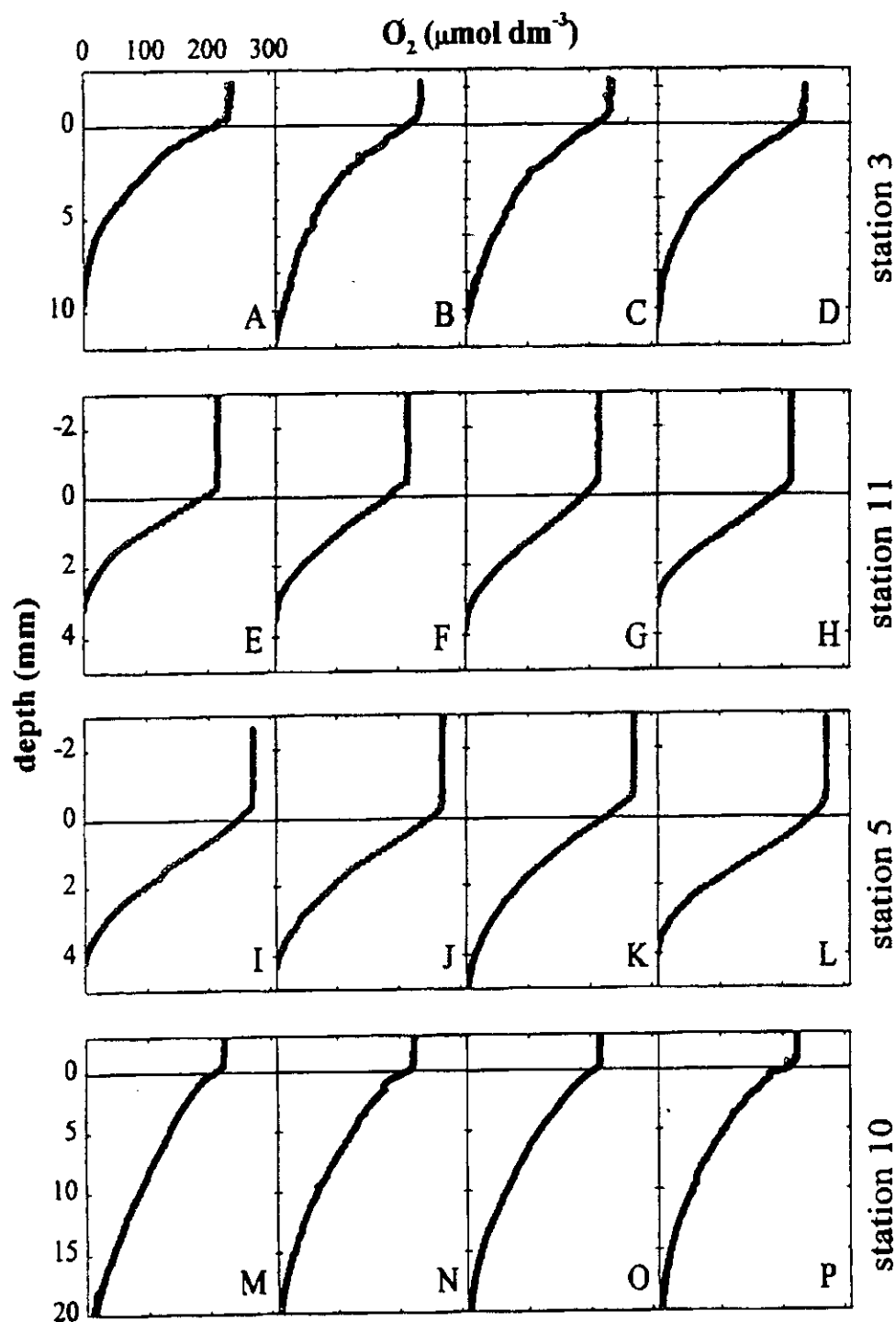


Fig. 6. *In situ* oxygen profiles and the best fit according to model 4. (A–D) station 3 in March 1992, (E–H) station 11 in August 1993, (I–L) station 5 in August 1993, and (M–P) station 10 in August 1993. Profiles from station 11 and 5 could only be described by assuming benthic photosynthesis.

Reactivity and Photosynthesis in Adriatic Sediments
(after Epping & Helder 1997)

$$0 = D_s \frac{d^2 C}{dz^2} - R$$

MODEL I:

$$\Sigma R(O_2)(z) = -R$$

Model II

$$\Sigma R(O_2)(z) = -R_0 \exp^{az}$$

Model III

Layer I

$$\Sigma R(O_2)(z) = -R_0$$

Layer II

$$\Sigma R(O_2)(z) = -R_0 e^{-a(z-z_1)}$$

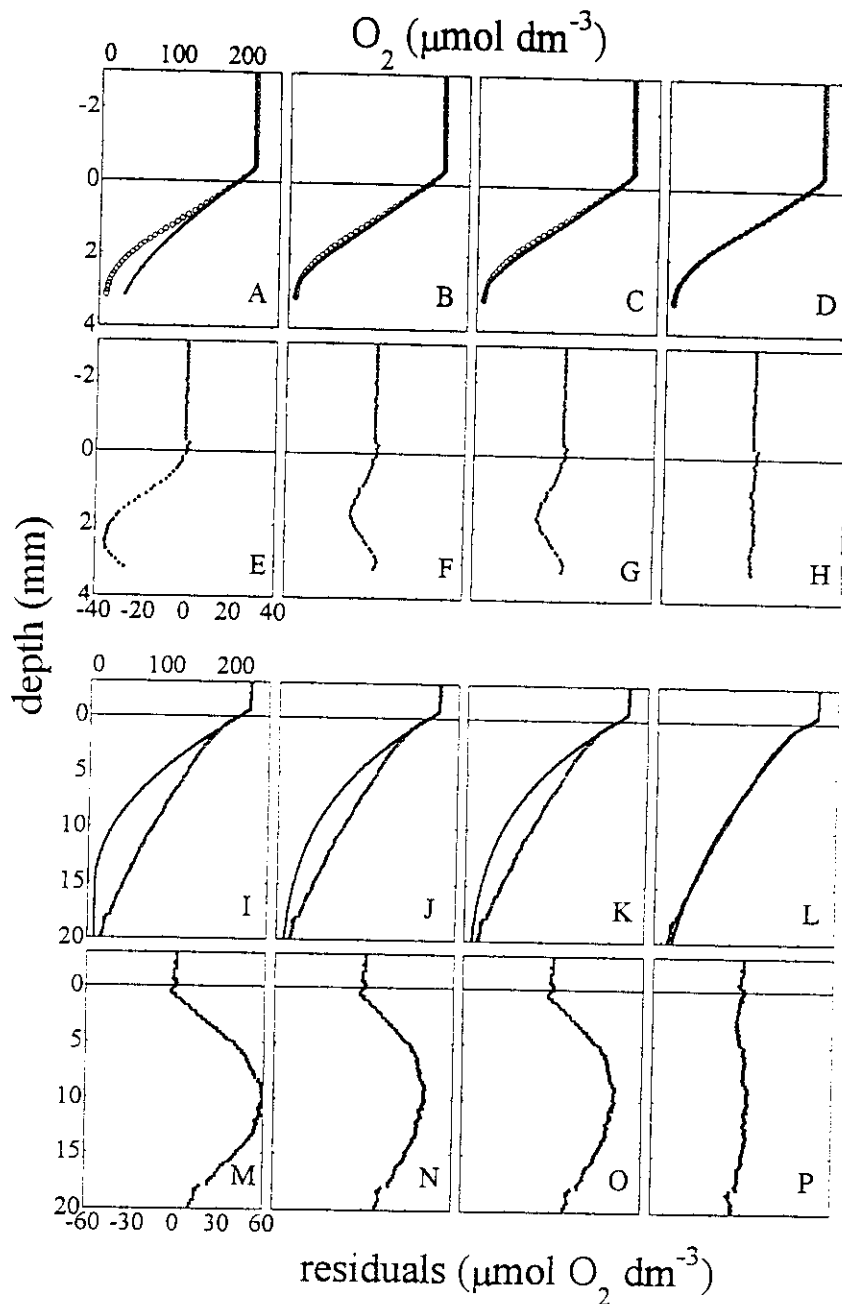
Model IV

Layer I

$$\Sigma R(O_2)(z) = -(R_1 + R_2)$$

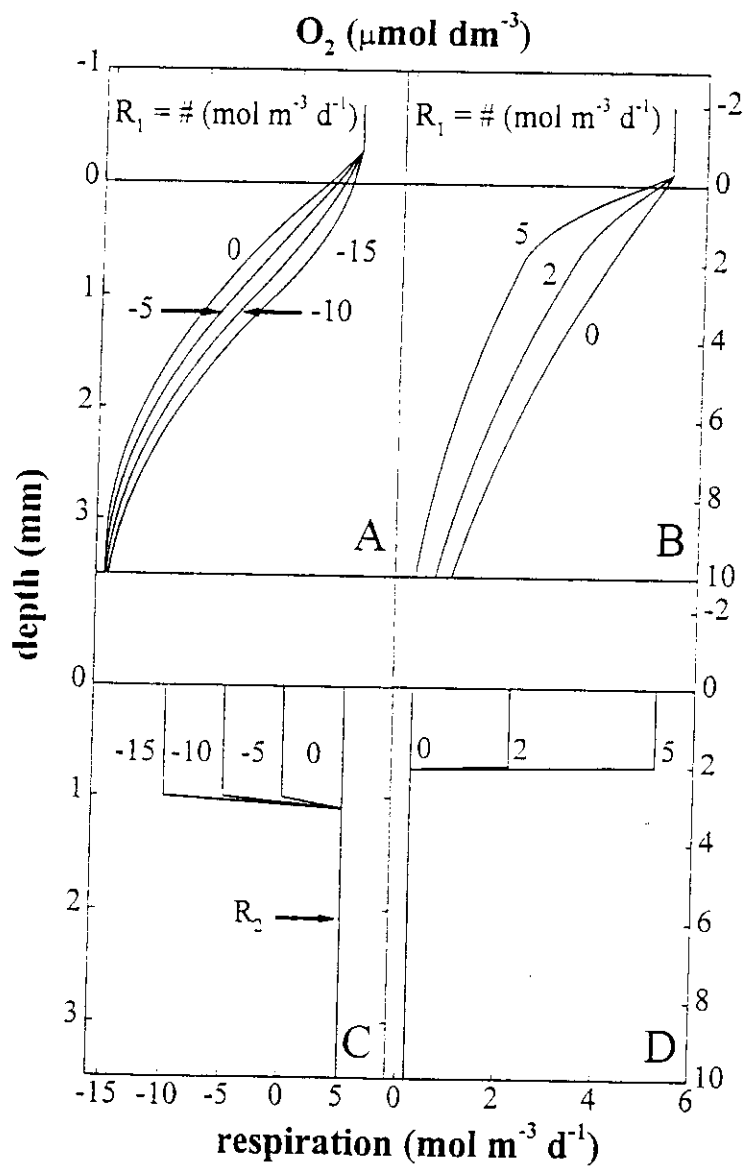
Layer II

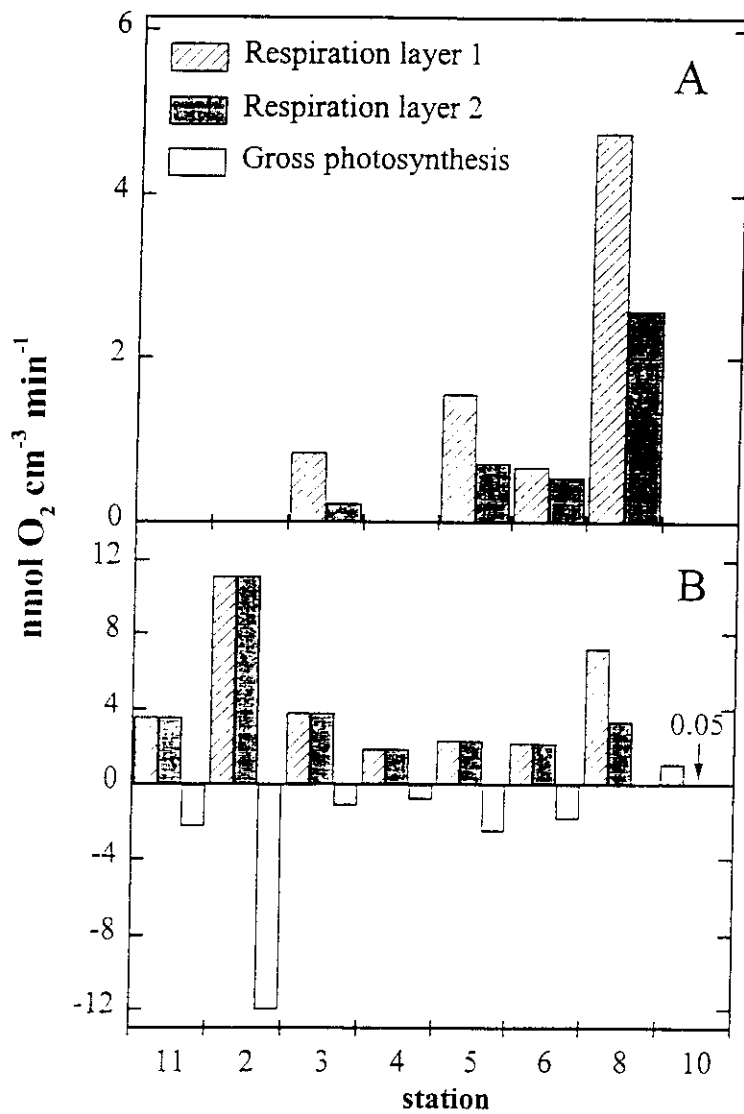
$$\Sigma R(O_2)(z) = -R_2$$



Epping and Helder. *Cont Shelf Res.* 17:1737-1764 (1998)

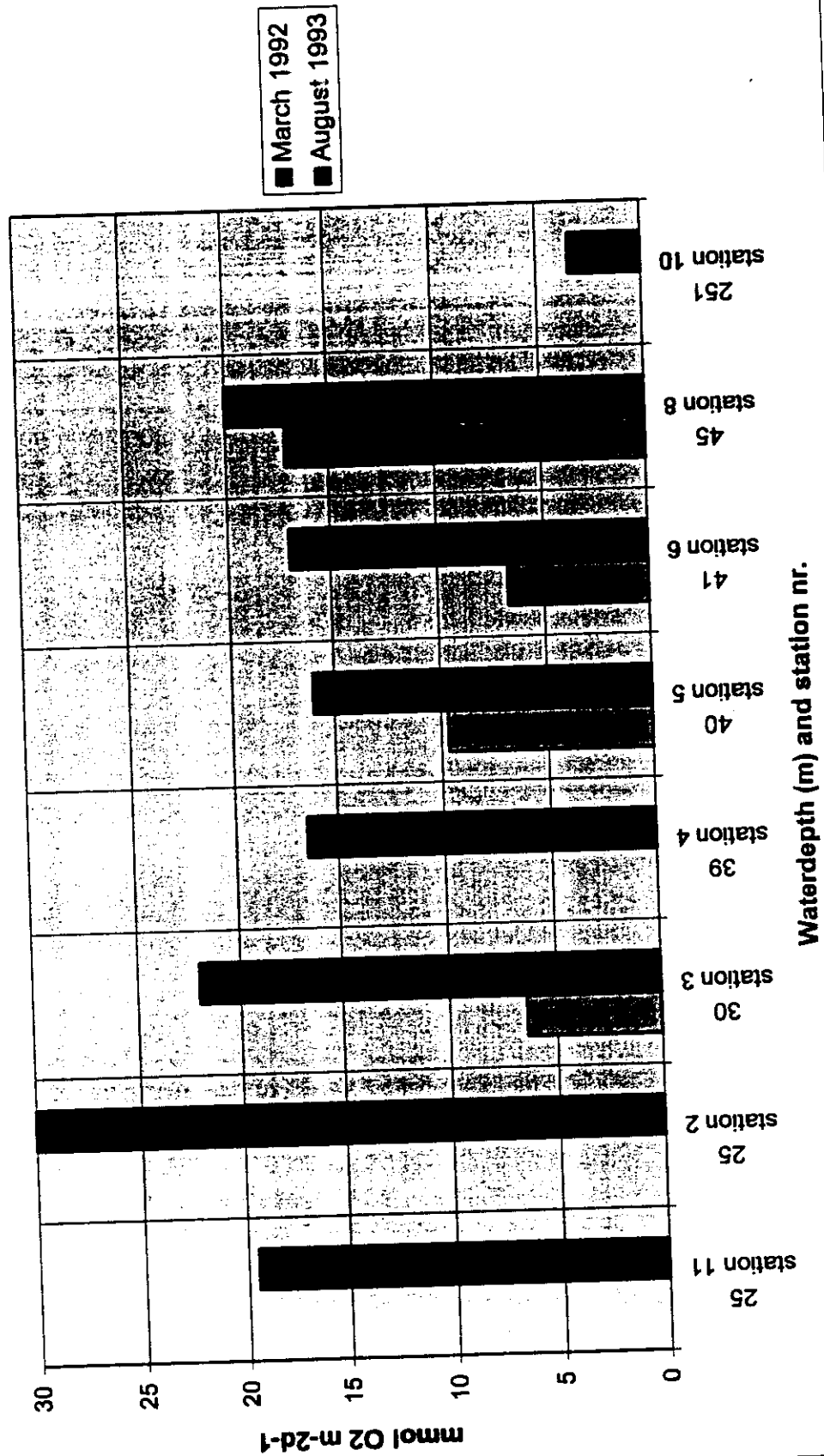
Figure 5. (A-L) Comparison of the four models (solid lines) with a representative in situ profile (symbols) for the shallow stations (A-D) and for the deep station (I-L). (E-P) Residuals, defined as the difference between the experimental and modeled value. (E-H) The residuals for the oxygen profile from the shallow station and (M-P) for the profile from the deep station. The left lane of panels refer to model i, the central left lane to model ii, the central right lane to model iii and the rightmost lane to model iv.





Epping + Kelders, 1998

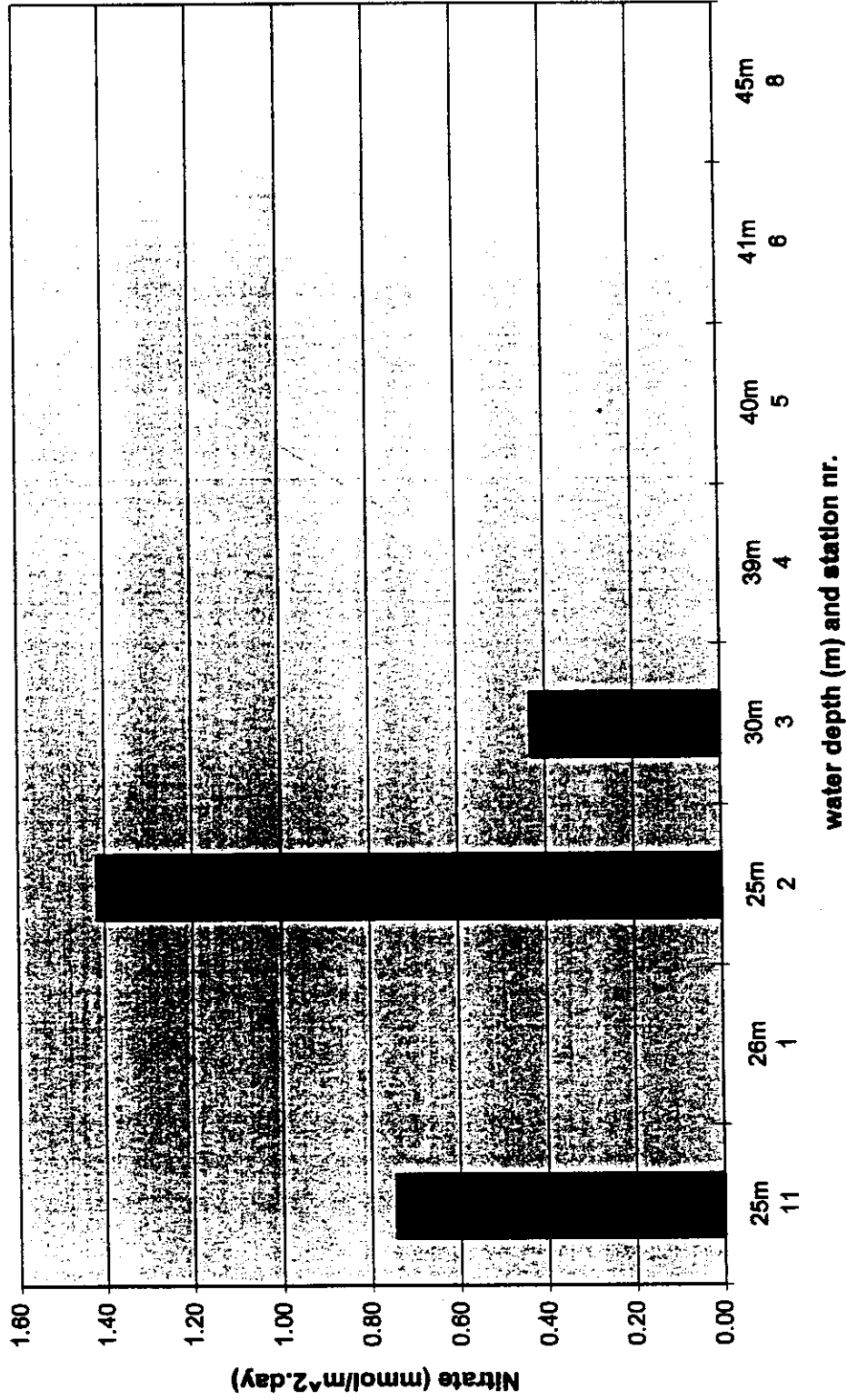
Benthic O2 fluxes



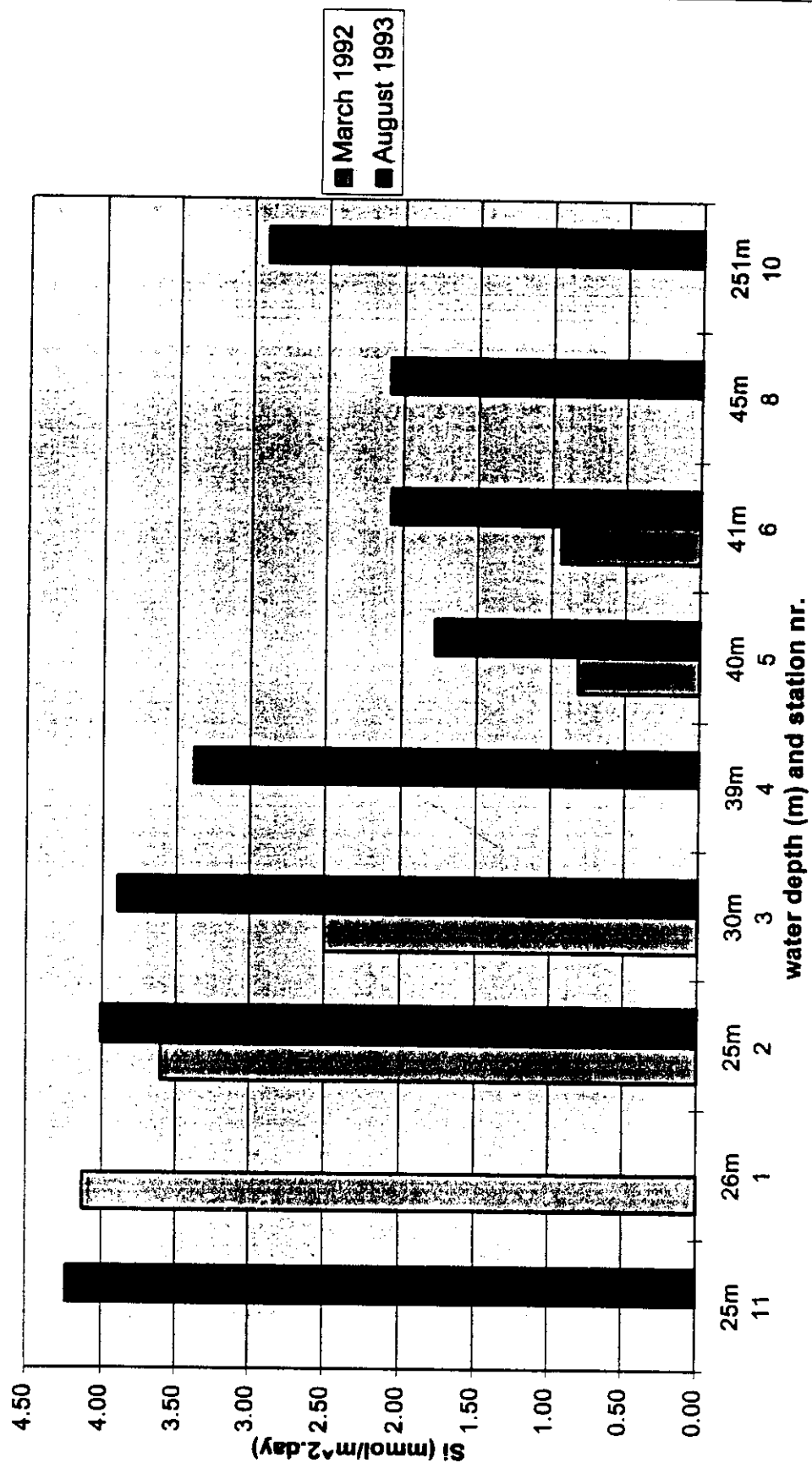
Benthic oxygen fluxes ($\text{mmol O}_2 \text{ m}^{-2} \text{ d}^{-1}$) in the shallow Northern Adriatic Sea

	Febr-March	August-Sept.
Giordani et al. (1992)	-	11.9 - 19.9
Tahey et al. (1996)	8.4 - 24.4	15.0 - 34.8
Epping and Helder (1997)	6.4 - 17.1	16.3 - 30.0
Moodley et al. (in press)	2.8 - 16.8	22.7 - 47.5

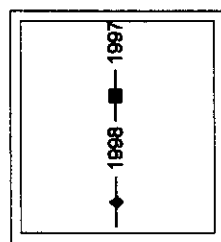
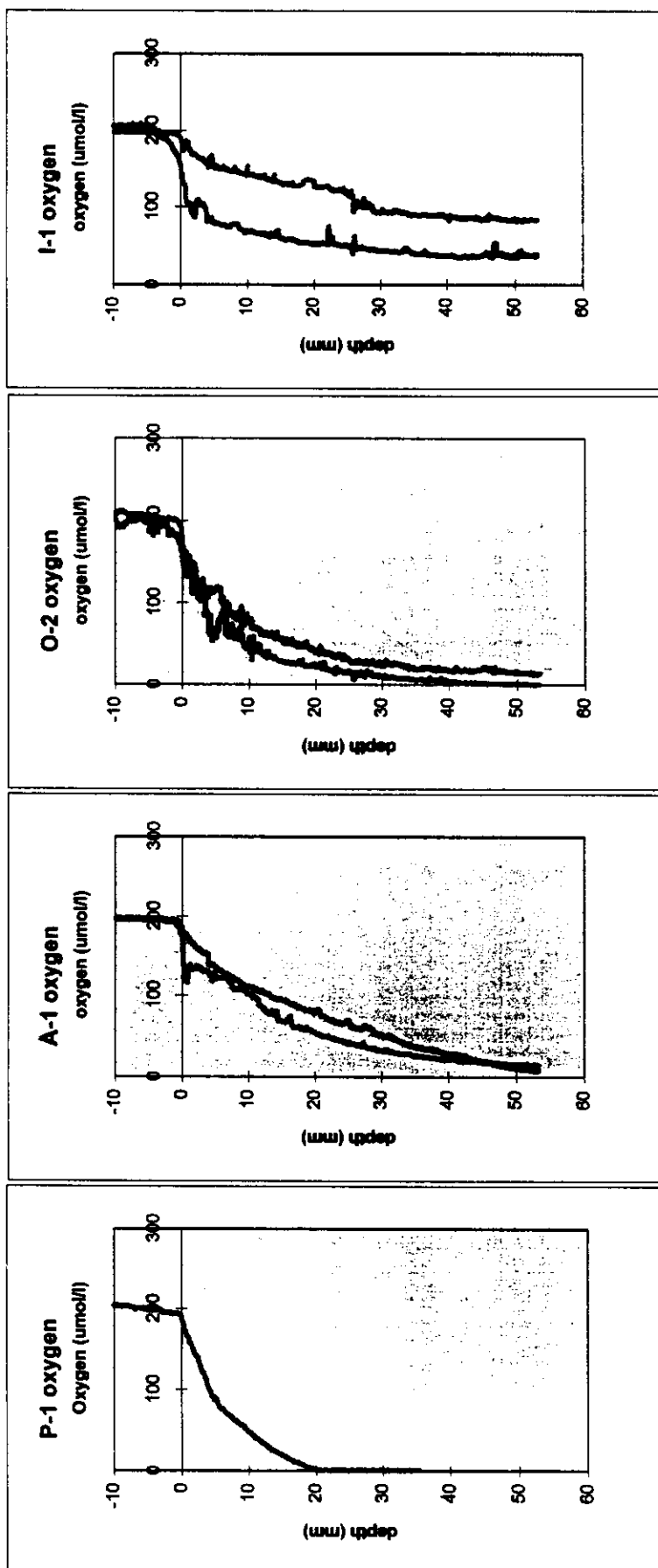
Benthic nitrate fluxes, STEP program

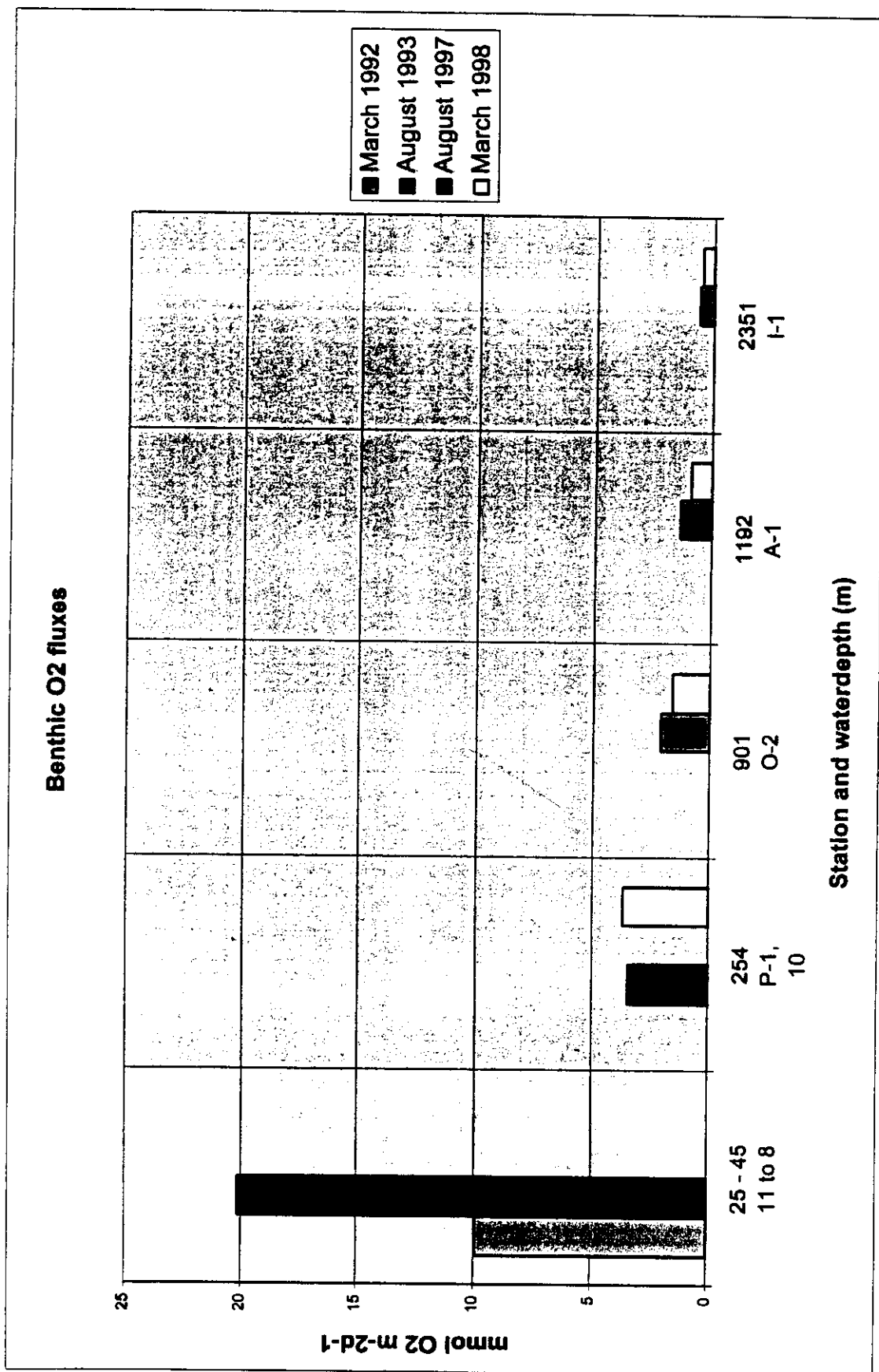


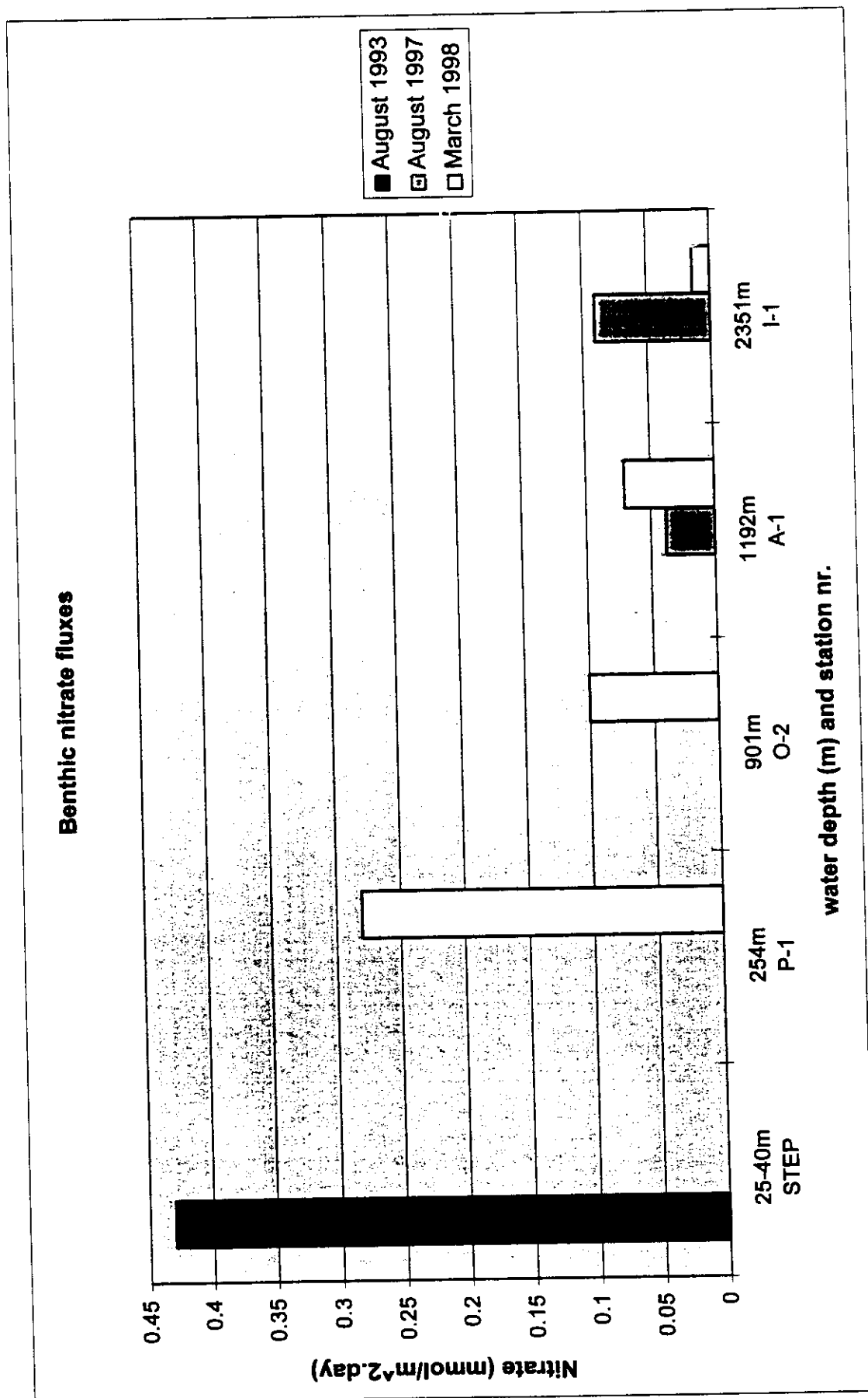
Benthic silica fluxes



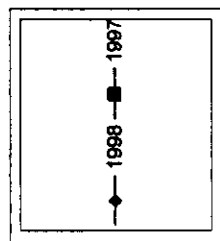
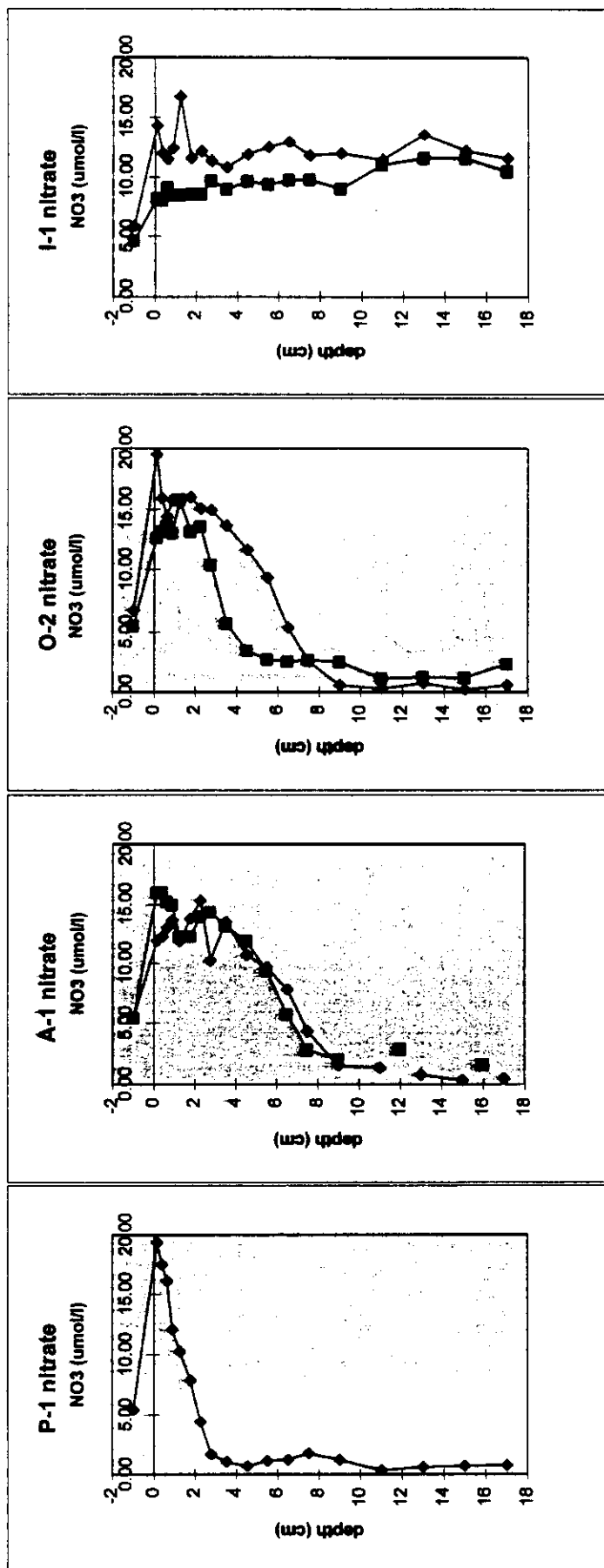
oxygen pore water profiles



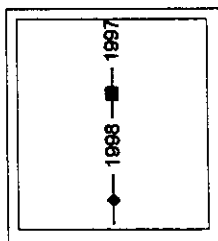
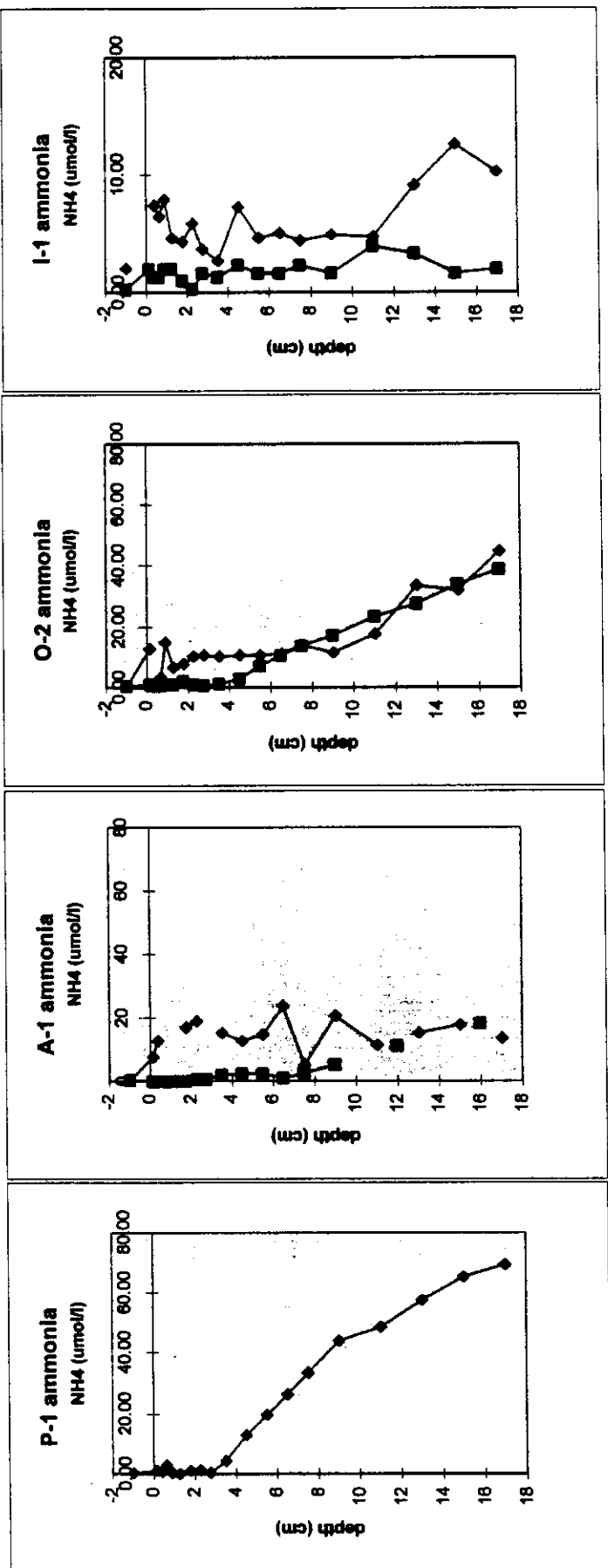




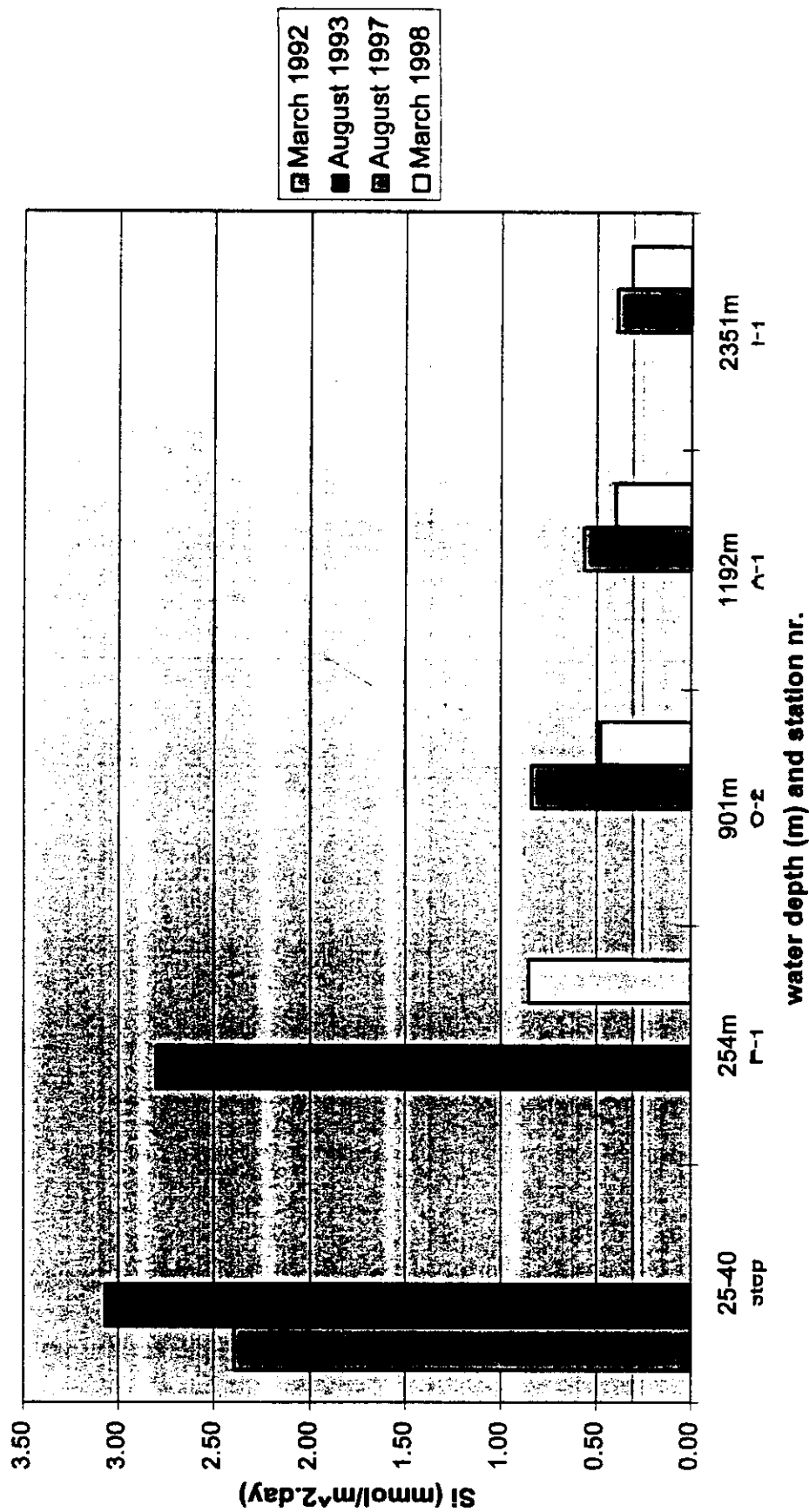
nitrate pore water profiles



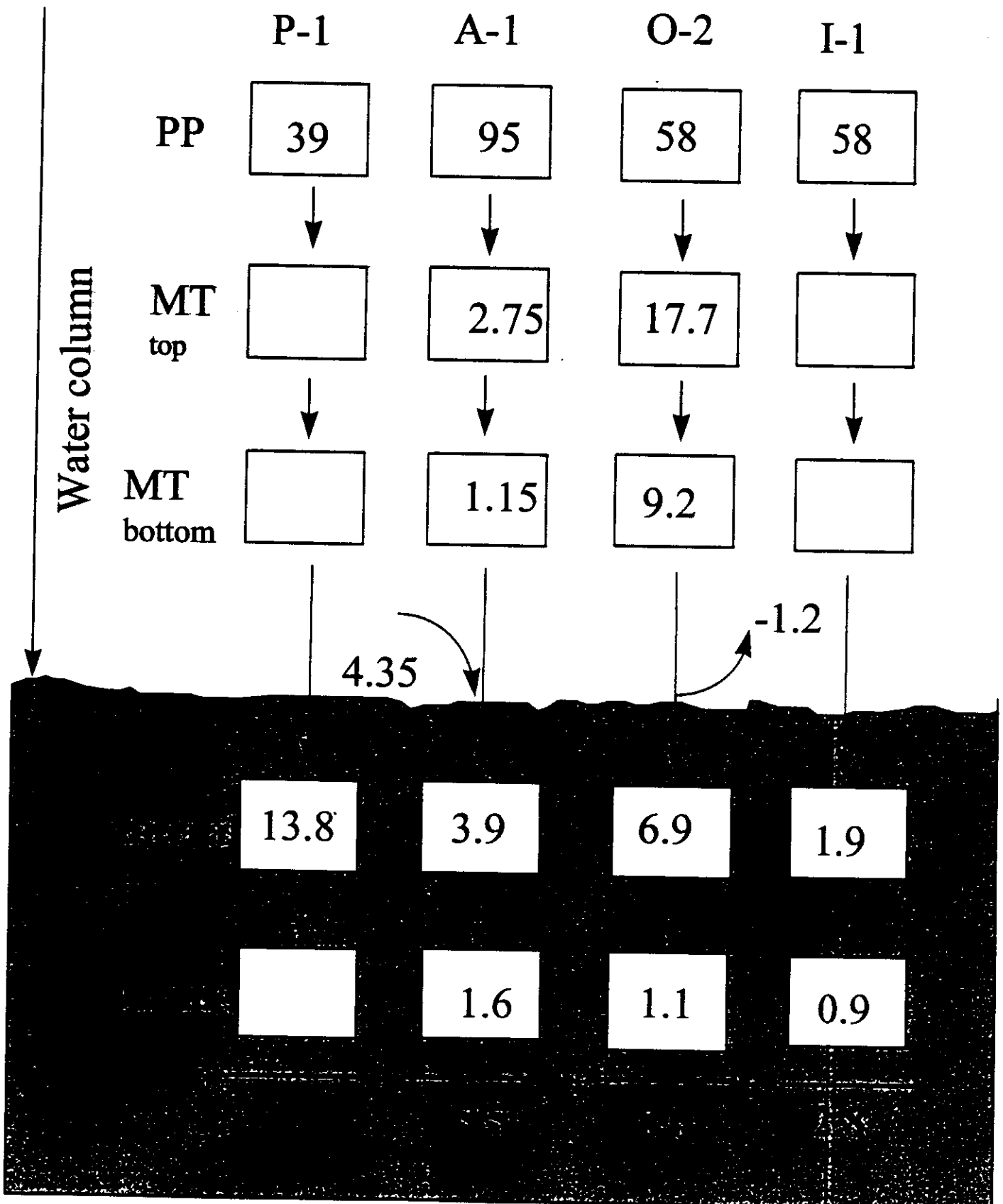
ammonia pore water profiles



Benthic silica fluxes

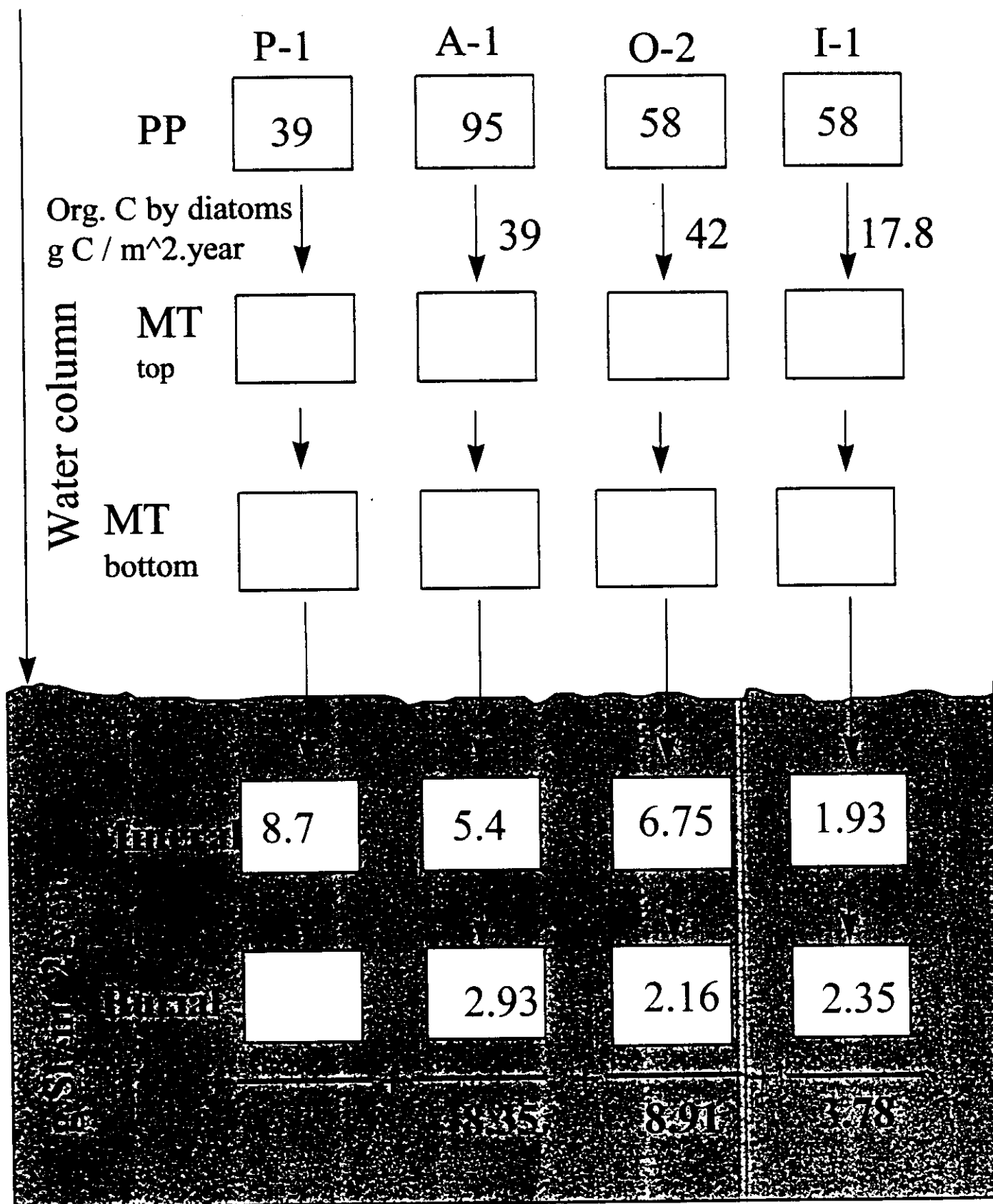


Organic Carbon (g.m-2.y-1)



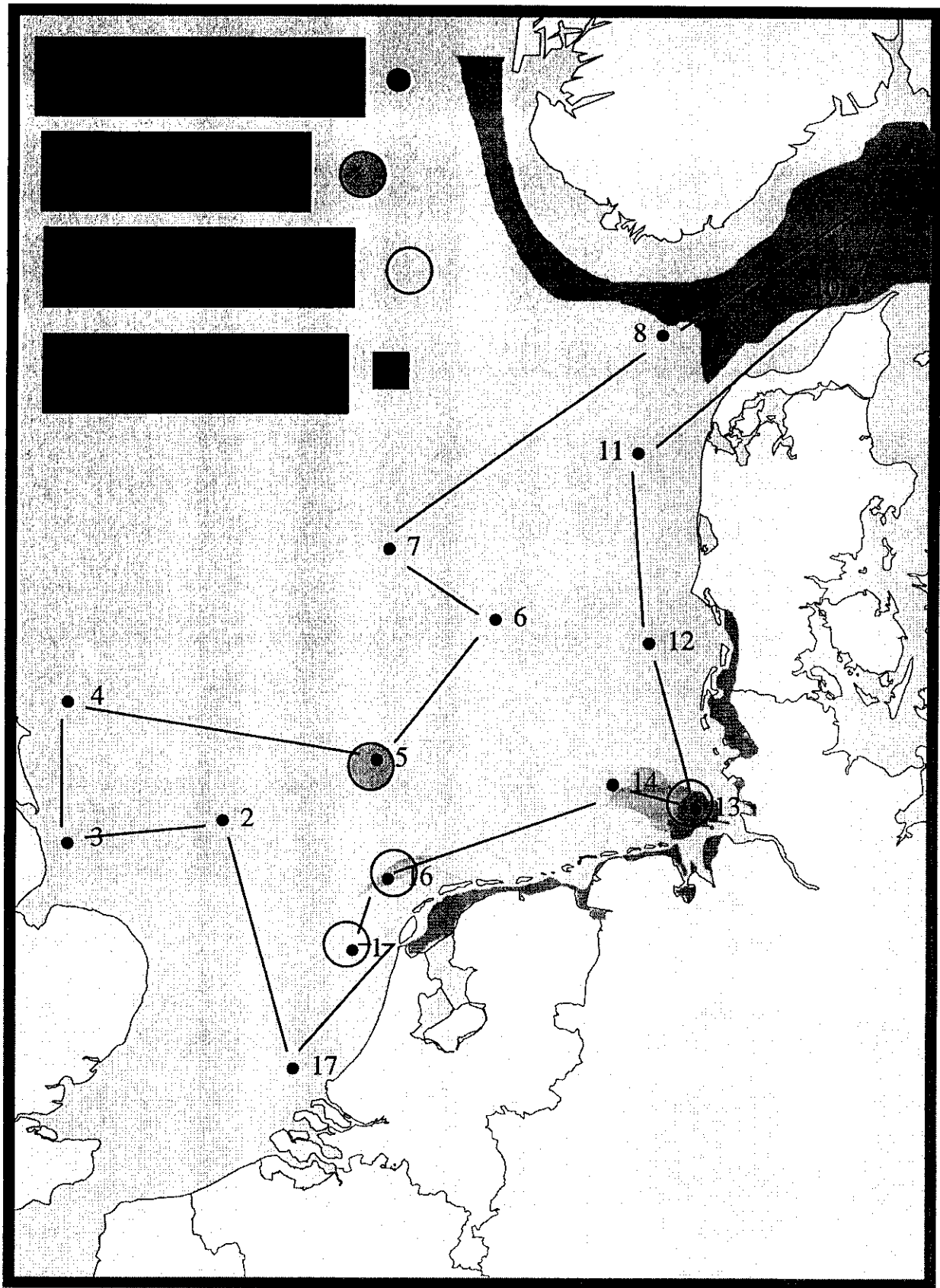
Contribution to org.C-budget by silicious organisms

Conversion of BioSi to org.C by $C/Si=4.7$ (w/w)

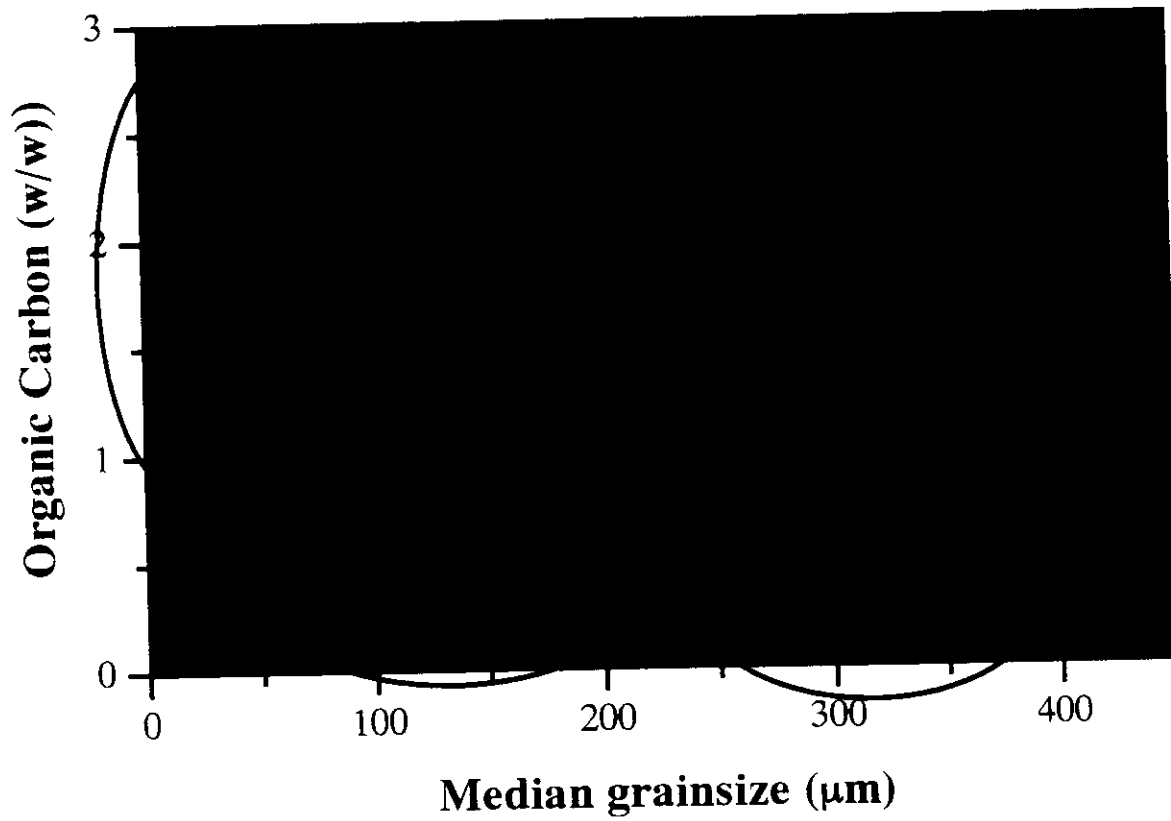


The Adriatic Sea: Conclusions

- *Benthic oxygen consumption in the Northern Adriatic shows a clear seasonal variation*
- *Oxygen budgets of these sediments indicate the occurrence of benthic primary production in Northern Adriatic sediments*
- *Benthic oxygen consumption rates accounts for 11 to 26% of the pelagic primary production in Winter. In summer, between 27 to 50% of the primary production is mineralised in the sediment, indicating the tight benthic-pelagic coupling*
- *In the mid-Adriatic depression, 15% of the primary production is mineralised in the sediment*
- *Sediments located in the southern Adriatic Sea account for 3 to 12 % of the pelagic primary production*
- *Benthic processes clearly affect the chemical composition of Adriatic water masses.*



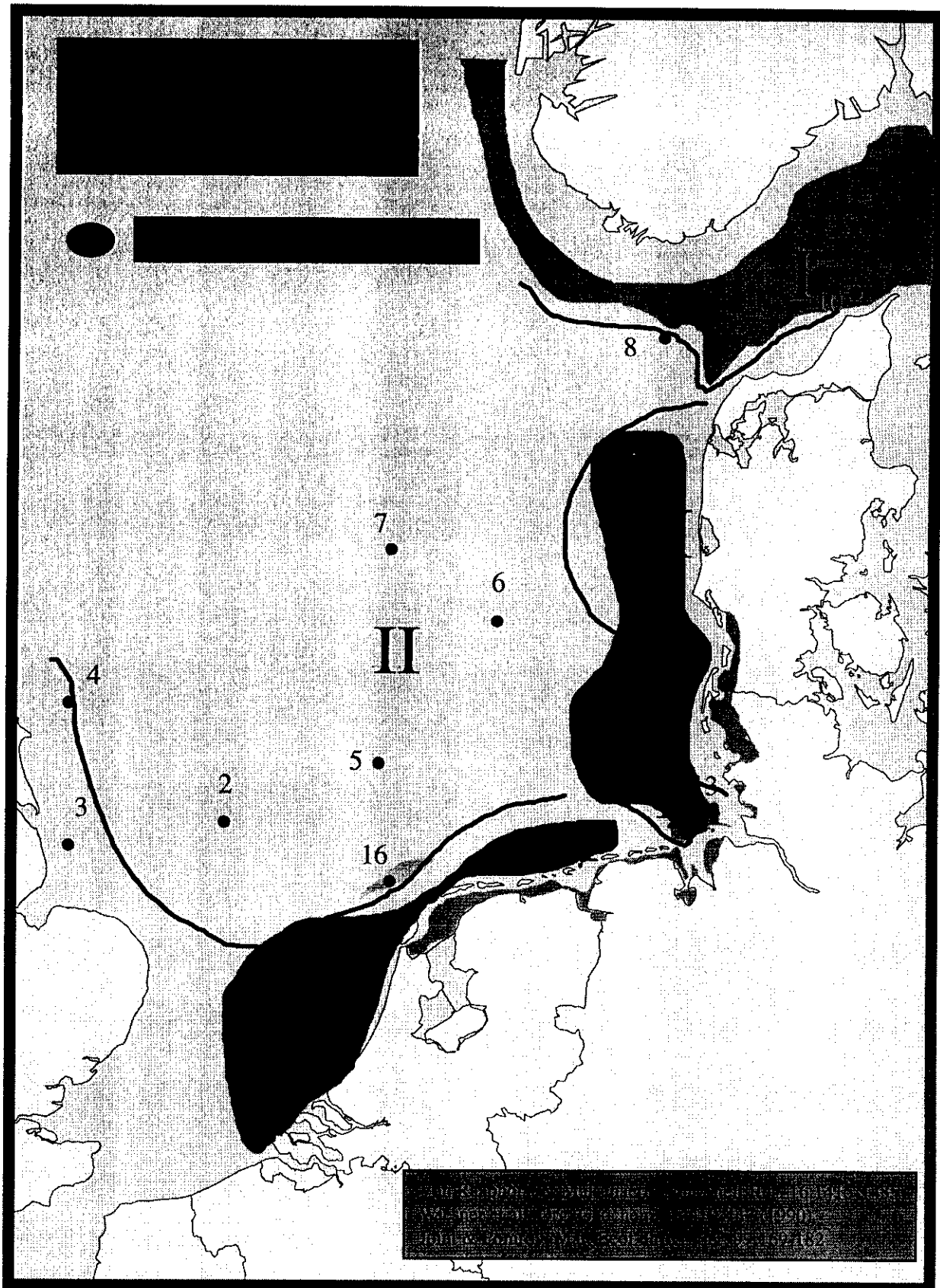
Grouping North Sea sediments

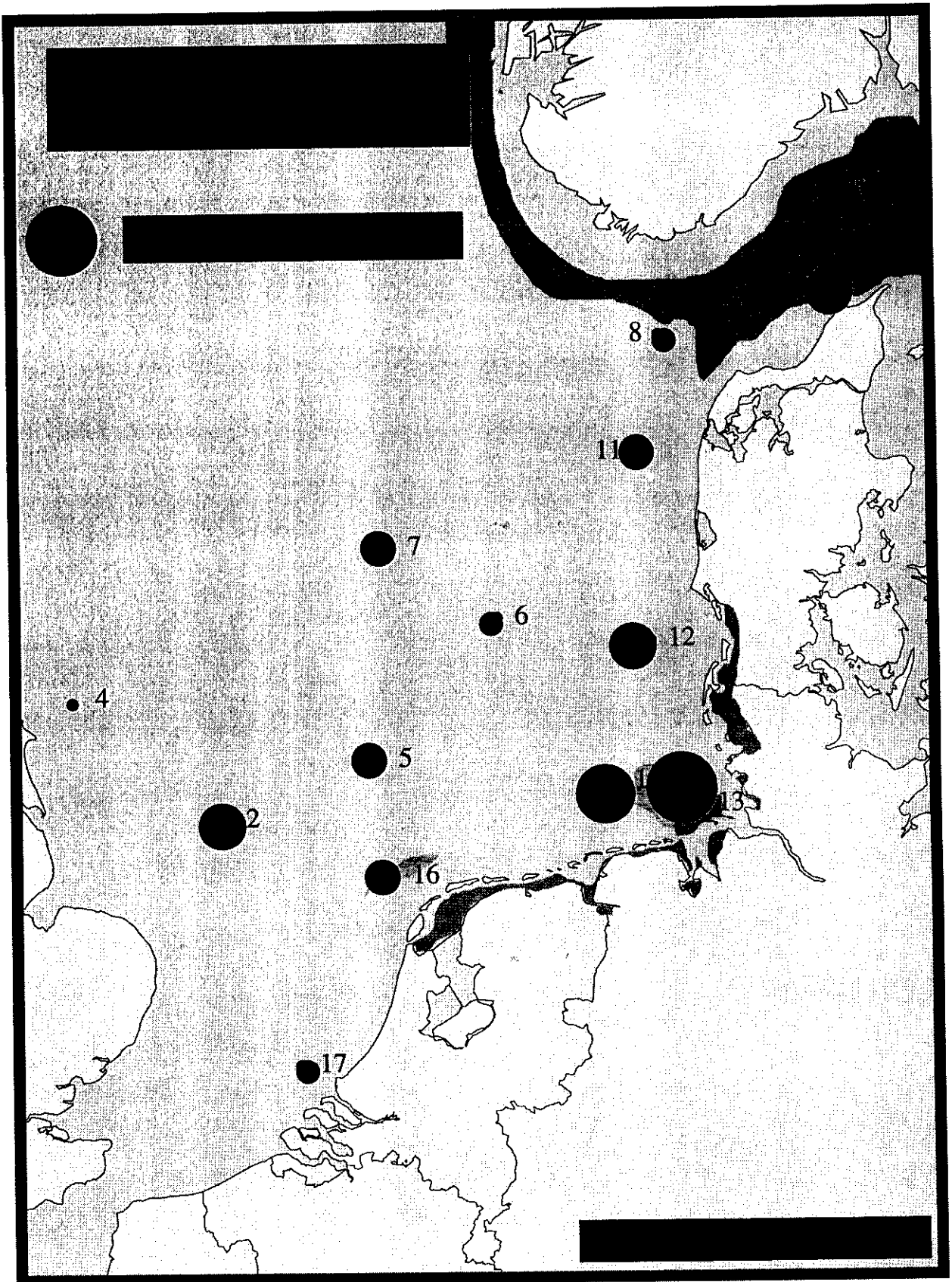


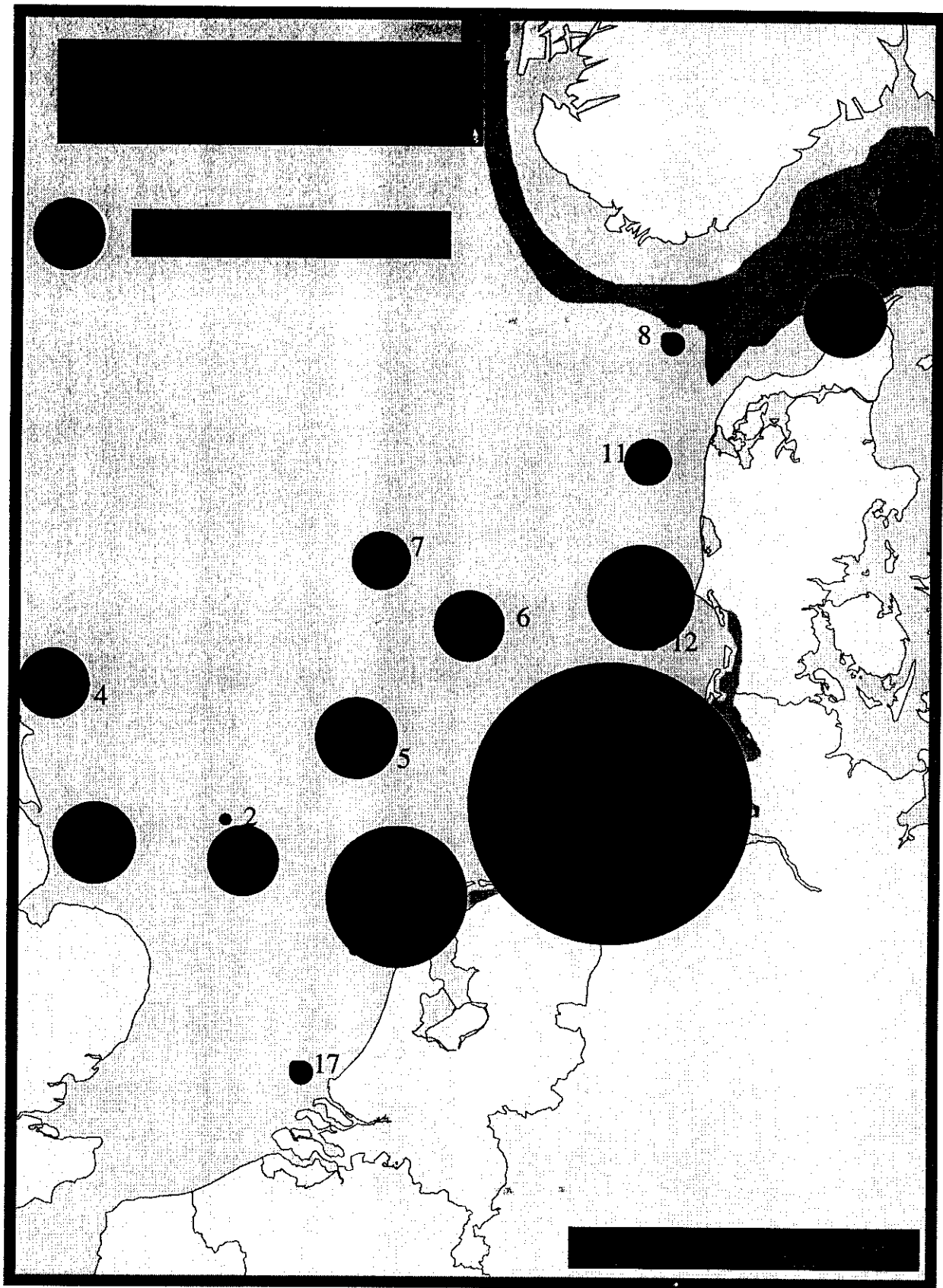
Lohse et al. *Ophelia* 42: 173-198 (1995)

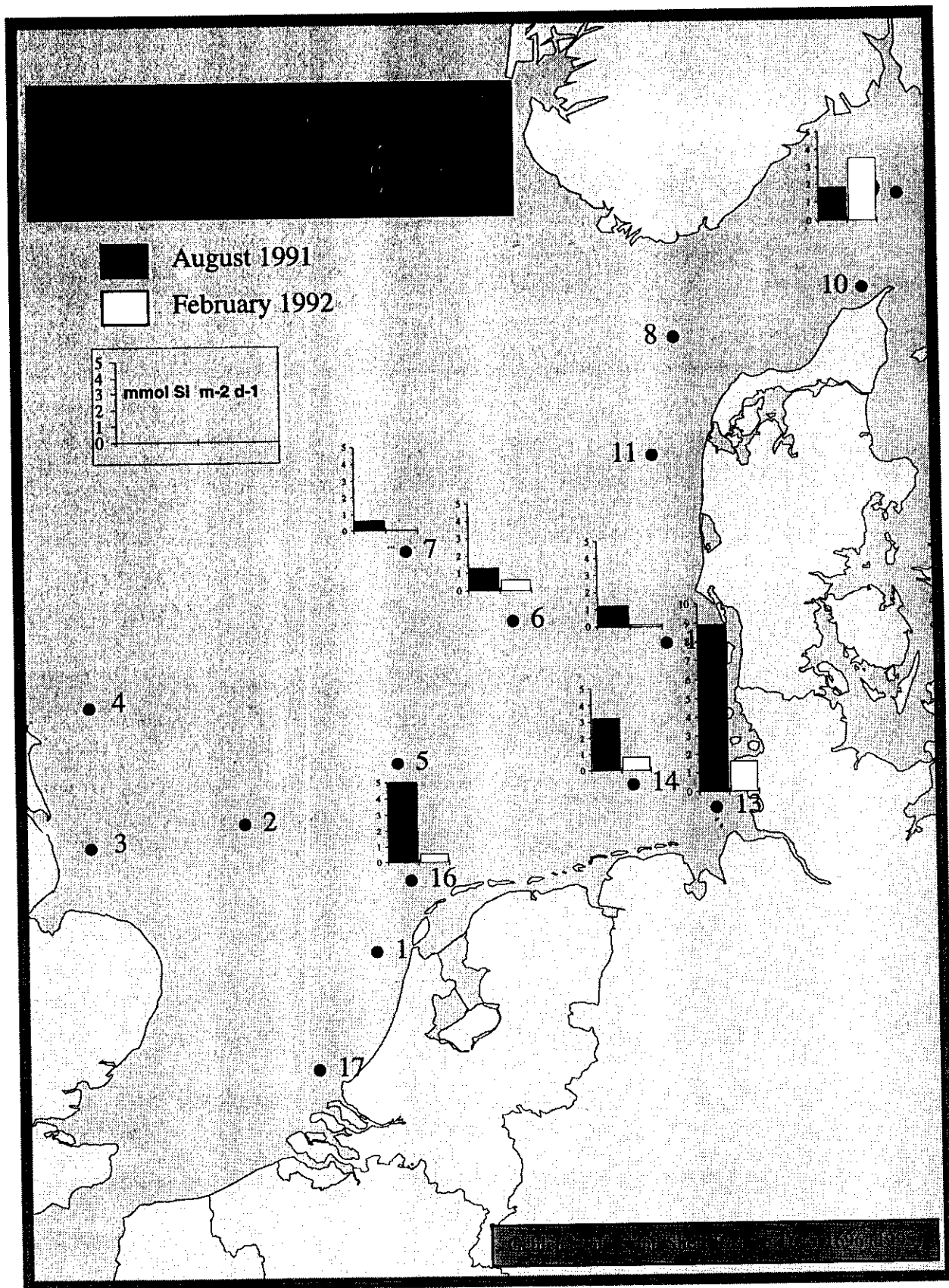


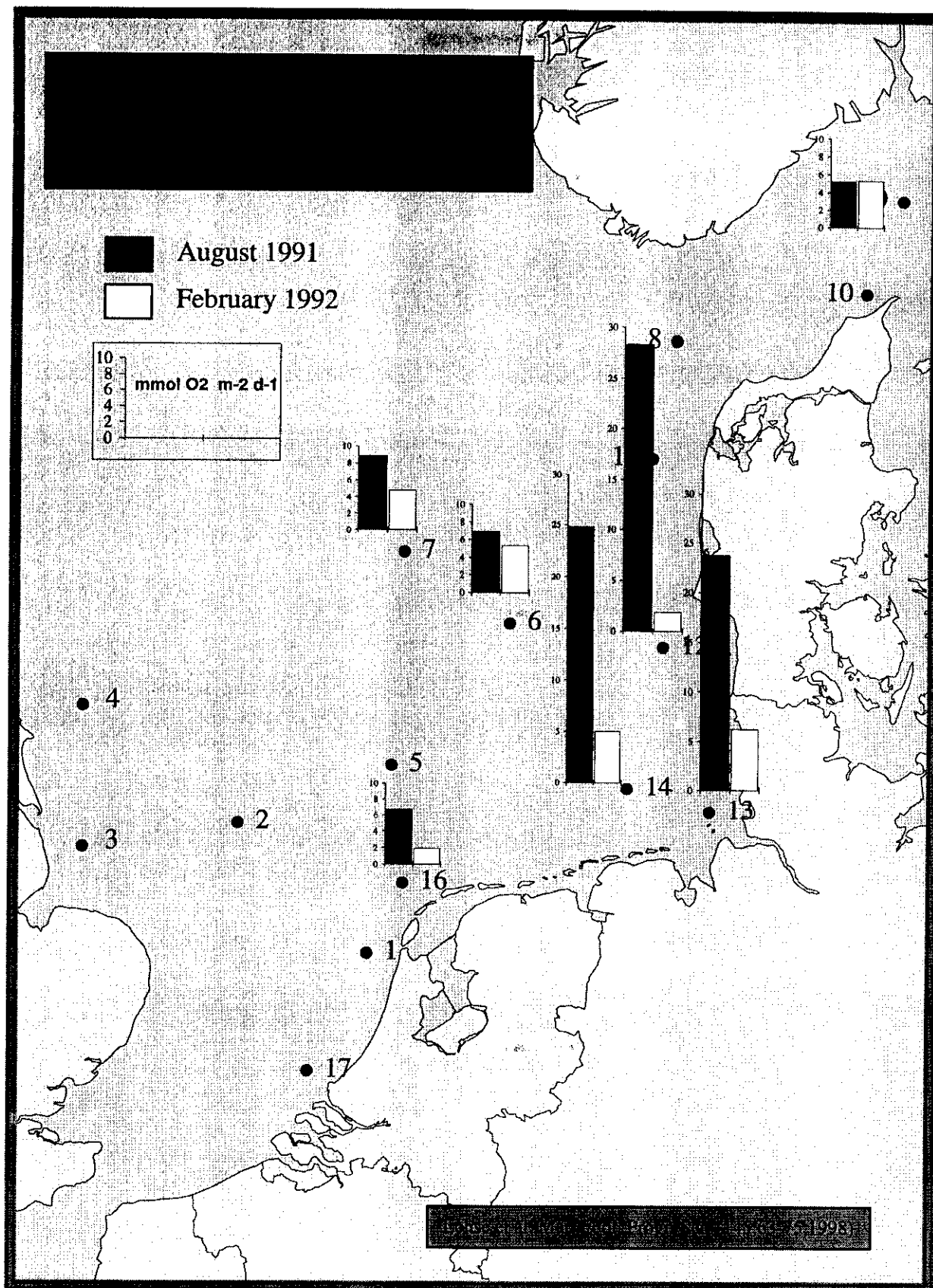
Oct. 96







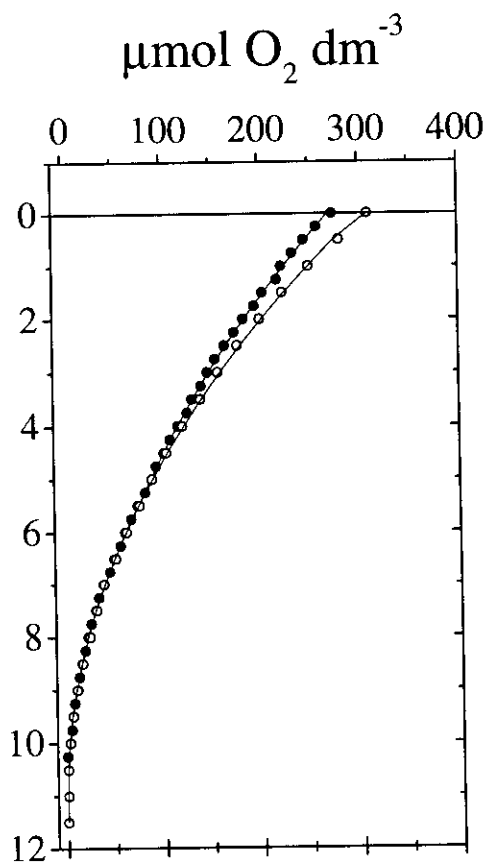




Skagerrak
water depth: 330 m

med. grainsize :
20 μm
(medium silt)

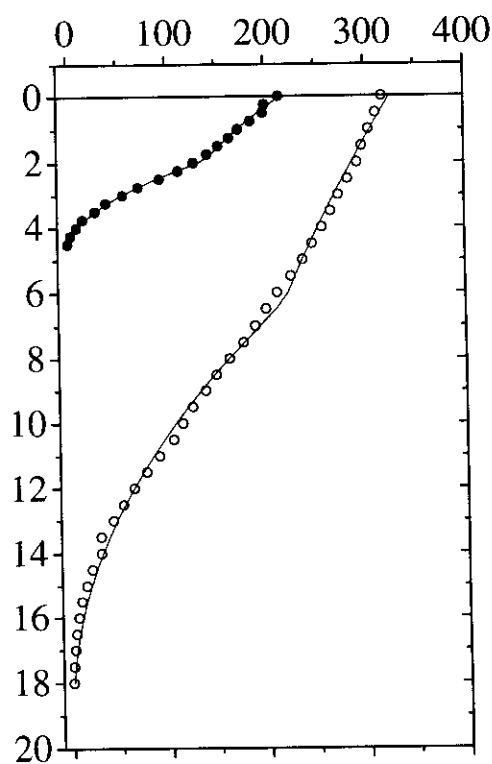
depth
(mm)



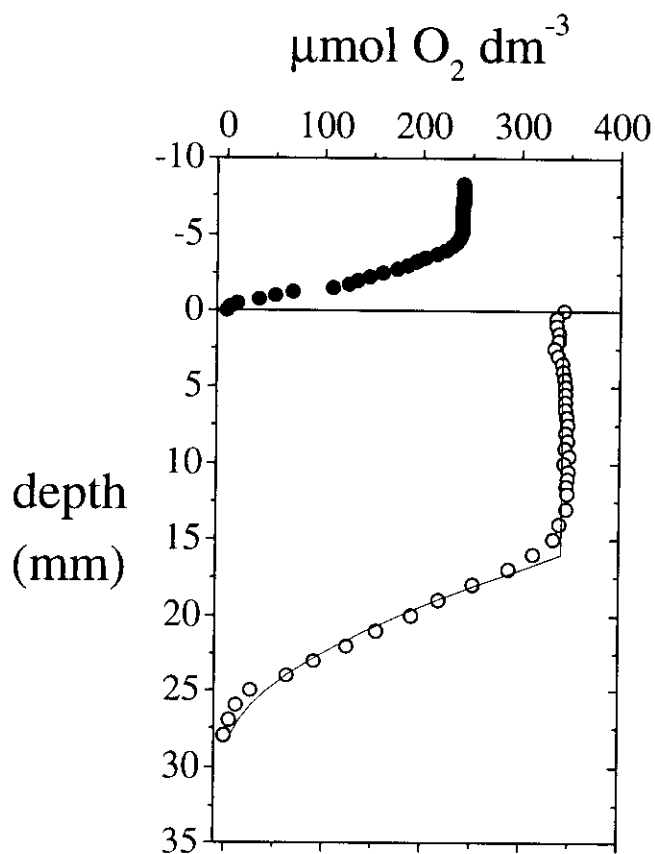
Frisian Front
water depth: 39 m

med. grainsize :
80 μm
(very fine sand)

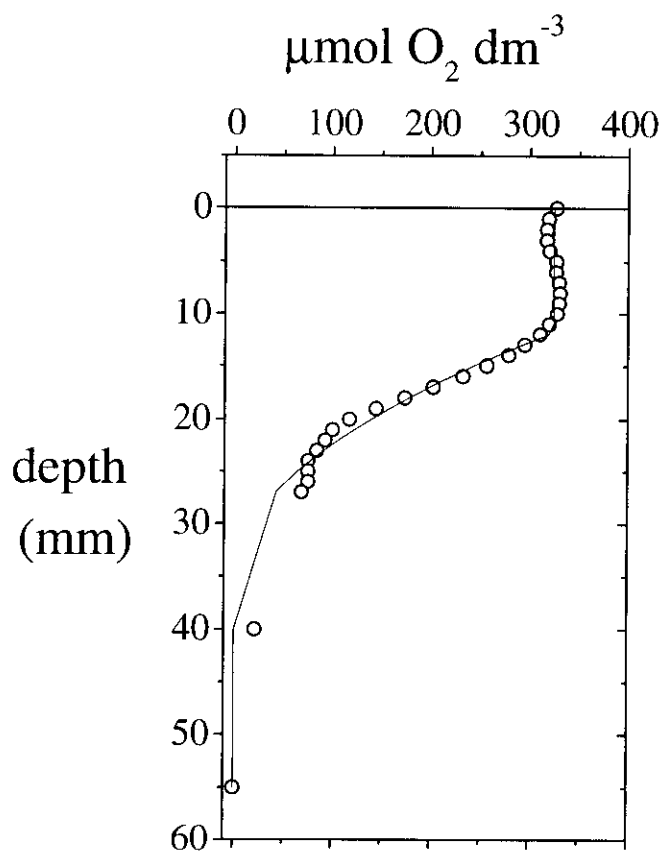
depth
(mm)

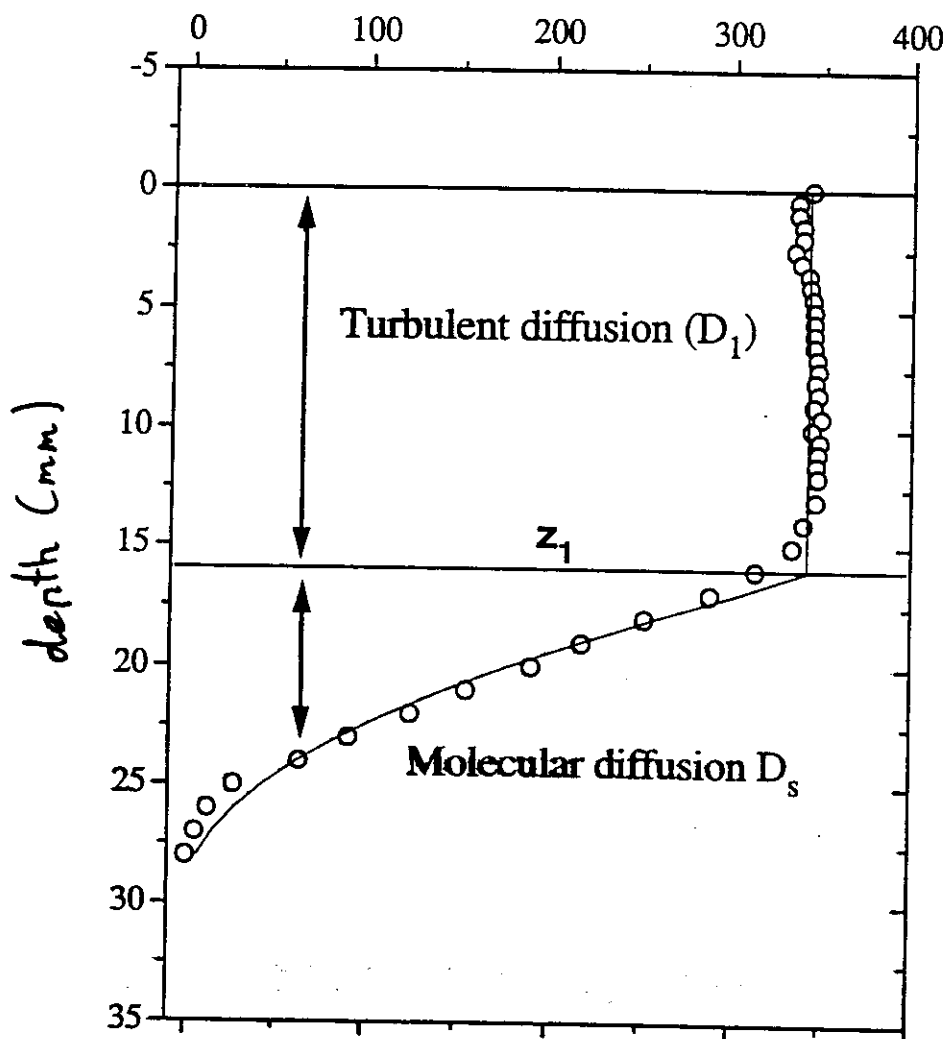


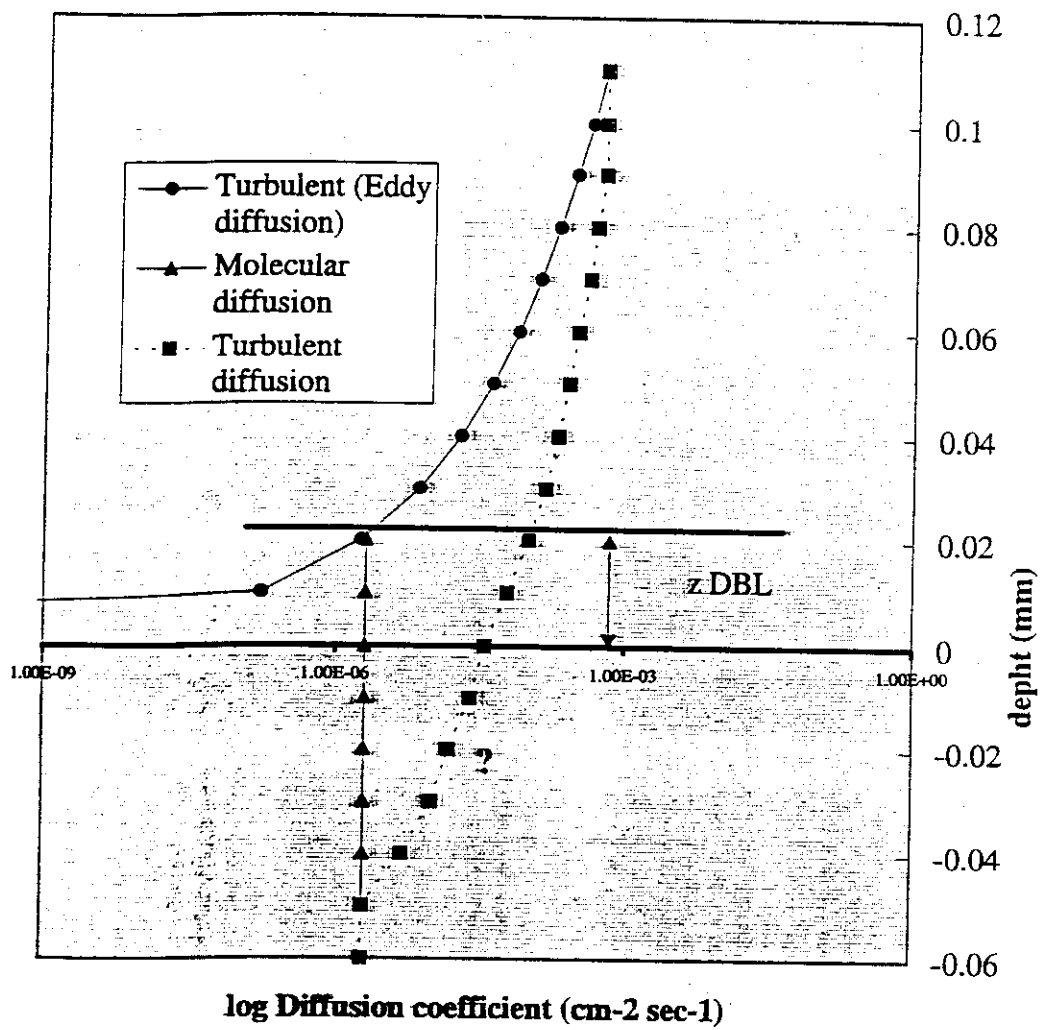
Danish coast
 water depth: 12 m
 medium grainsize:
 190 μm
 (fine sand)

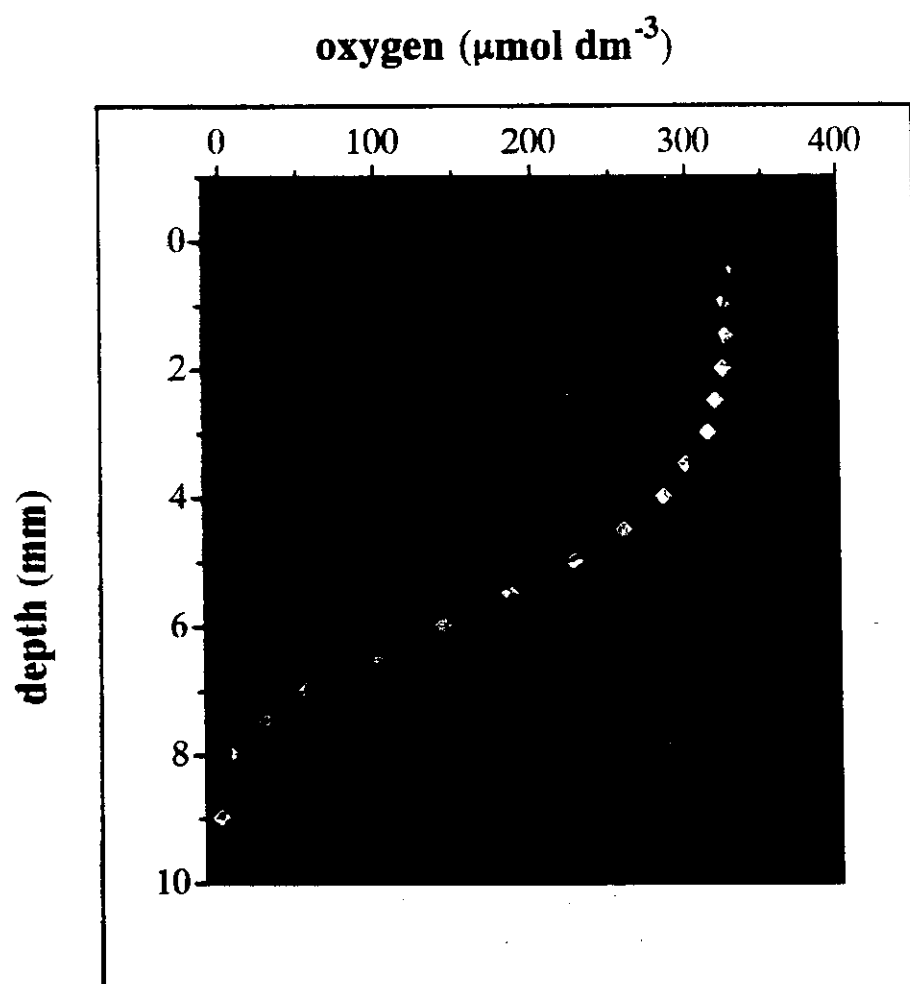


Doggerbank
 water depth: 60 m
 medium grainsize:
 300 μm
 (medium sand)









- ♦ 10 minutes
- ▲ 20 minutes
- 25 minutes
- 30 minutes

after box core retrieval

Time scales for molecular and turbulent diffusion

$$t = z^2 / 2D_z$$

at $T = 20^\circ\text{C}$, $D = 32 \times 10^{-6} \text{ cm}^2/\text{sec}$, $\phi = 0.2$

Distance (z) (cm)	Molecular diffusion coefficient	Turbulent diffusion coefficient
coefficient	6.4	429
(x 10 ⁻⁶ cm ² -sec ⁻¹)		
0.01	7.8 sec	0.12 sec
0.02	3 min	2 sec
0.1	13 min	11 sec
0.2	2 h	2 min
1	21 h	19 min
2	23 d	8 h

Conclusions

- *Near bottom pressure fluctuations caused by tidal currents and wave action may induce turbulent transport processes in "non-depositional" continental shelf sediments.*
- *Oxic/anoxic oscillations may lead to a more complete mineralisation of organic matter*

Further studies

- *Lander technologies to study the processes under in-situ conditions*
- *Calibration of stirring regimes in incubation chambers*
- *Laboratory experiments (thin-plug techniques)*



Dec. 96



Conclusions

North Sea sediments act as link rather than sink.

Temporal deposition of relatively fresh material causes large seasonal differences of biogeochemical processes in mid-shelf sediments. These sites and particularly those with more constant deposition (e.g. German Bight) are key areas for benthic mineralisation.

Before reaching the final depocentre in the Skagerrak, most of the organic matter has been decomposed. The material deposited and buried here is largely refractory.



Adriatic vs North Sea Biogeochemical characteristics of depositional areas

<i>Summer</i>	<i>Adriatic (Po Plume)</i>	<i>North Sea (German Bight)</i>
---------------	--------------------------------	---

Primary production ($\text{mmol C m}^{-2} \text{ d}^{-1}$)	50	110
Sedimentation rate (cm yr^{-1})	0.24-0.29	0.5-0.8
Sediment composition	clay-silt	medium silt
Oxygen uptake ($\text{mmol O}_2 \text{ m}^{-2} \text{ d}^{-1}$)	15-25	50-70
Nitrogen mineralisation ($\text{mmol N m}^{-2} \text{ d}^{-1}$)	2-3	12
% primary production recycled in sediments	40	80
burial efficiency (%)	50	20

Adriatic vs North Sea Biogeochemical characteristics of non-depositional areas

<i>Summer</i>	<i>Adriatic (>30m)</i>	<i>southern North Sea</i>
---------------	-------------------------------	-------------------------------

Primary production ($\text{mmol C m}^{-2} \text{ d}^{-1}$)	40	26
Sedimentation rate (cm yr^{-1})	0	0
Sediment composition	medium sand	medium sand
Oxygen uptake ($\text{mmol O}_2 \text{ m}^{-2} \text{ d}^{-1}$)	5	10-15
Nitrogen mineralisation ($\text{mmol N m}^{-2} \text{ d}^{-1}$)	0.2	0.5
% primary production recycled in sediments	20	30-50
burial efficiency (%)	0	0

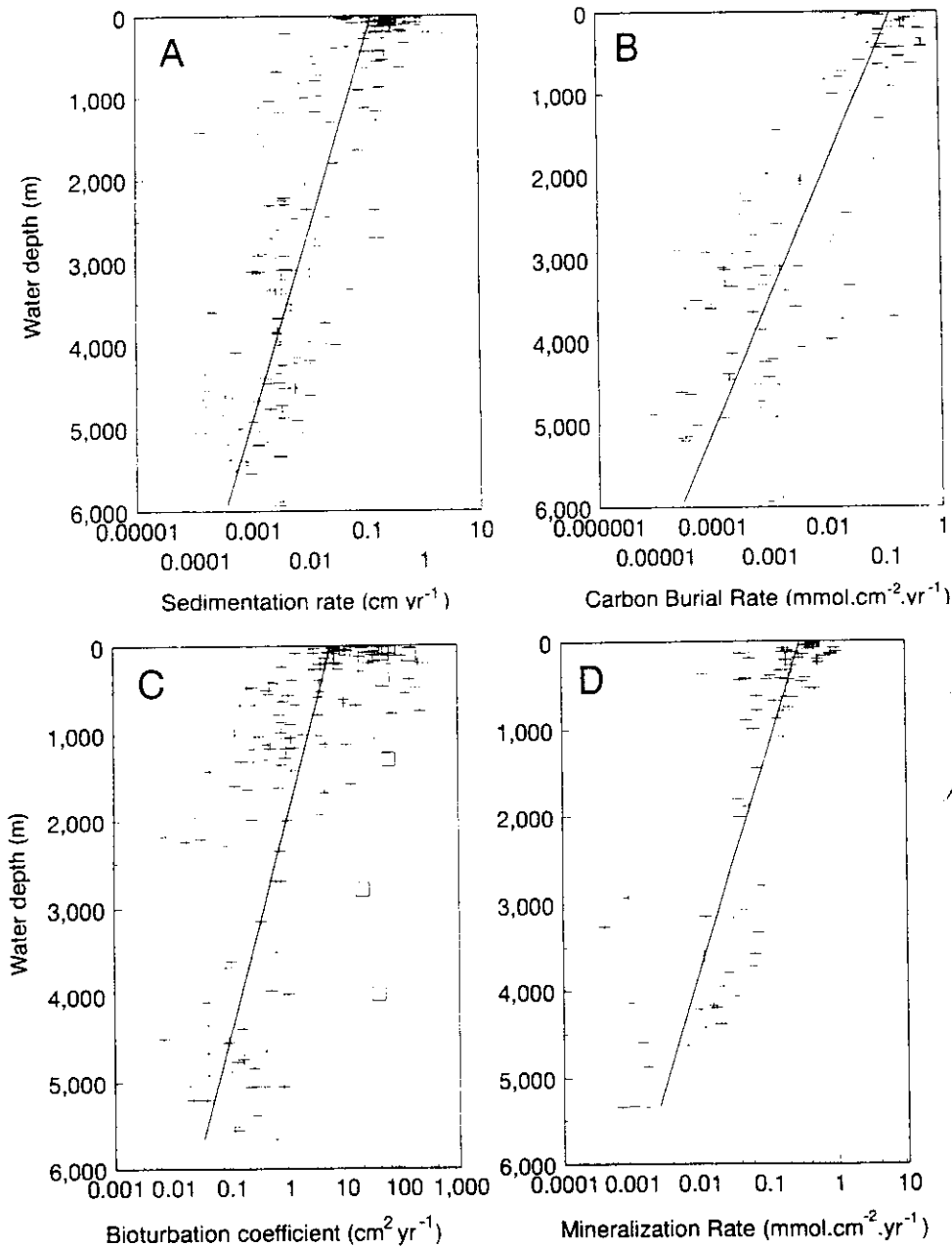


Fig. 3. Parameter versus water depth (m) plots. The regression curves based on the coefficients listed in Table 3 are included as well (solid lines).

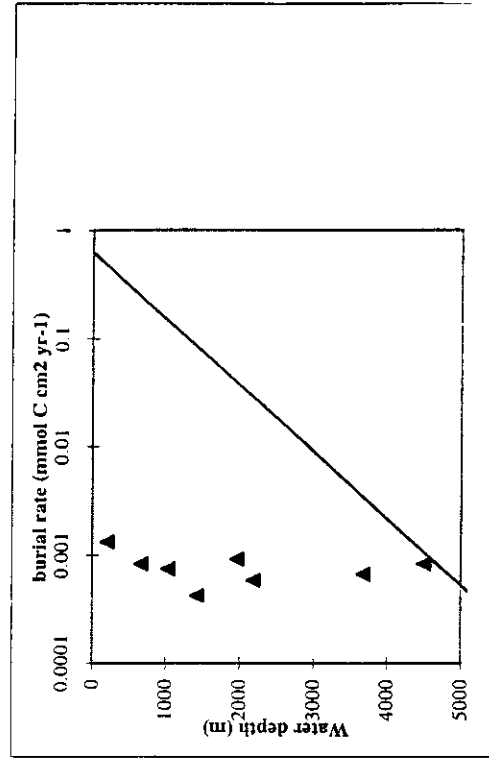
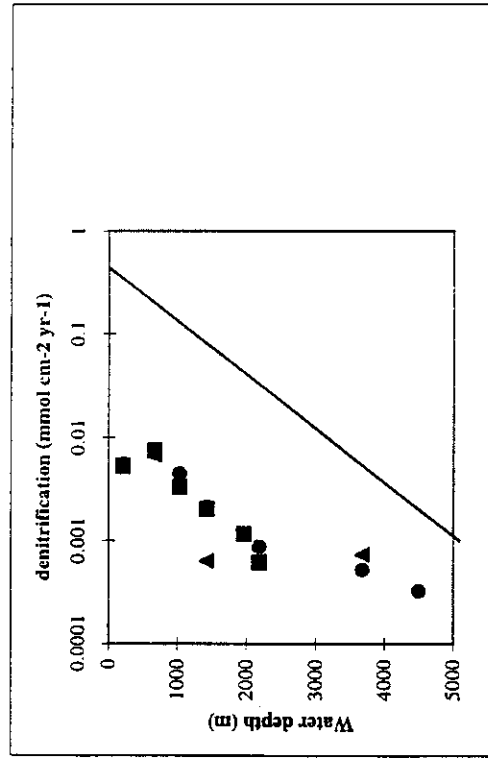
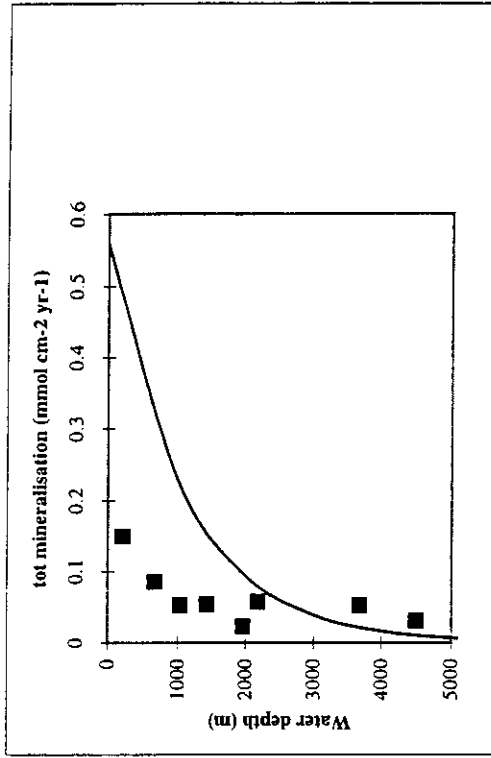
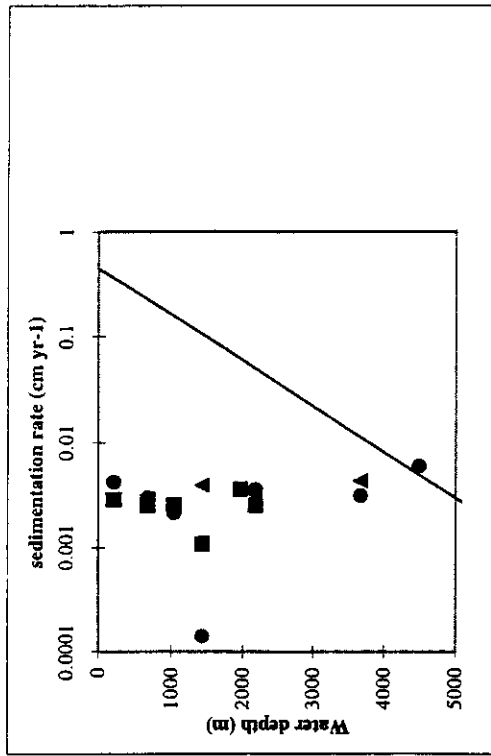
- (A) Sediment accumulation rate (ω ; cm year^{-1}).
 (B) Organic carbon burial rate (F_b ; $\text{mmol C cm}^{-2} \text{ year}^{-1}$).
 (C) $D_b\text{-}^{210}\text{Pb}$ (crosses; $\text{cm}^2 \text{ year}^{-1}$) and $D_b\text{-}^{234}\text{Th}$ (open squares; $\text{cm}^2 \text{ year}^{-1}$).
 (D) Total mineralization rate (F_o ; $\text{mmol C cm}^{-2} \text{ year}^{-1}$).

McCallum et al
1997
J. exp. Geol. Res.
44: 32 = 344

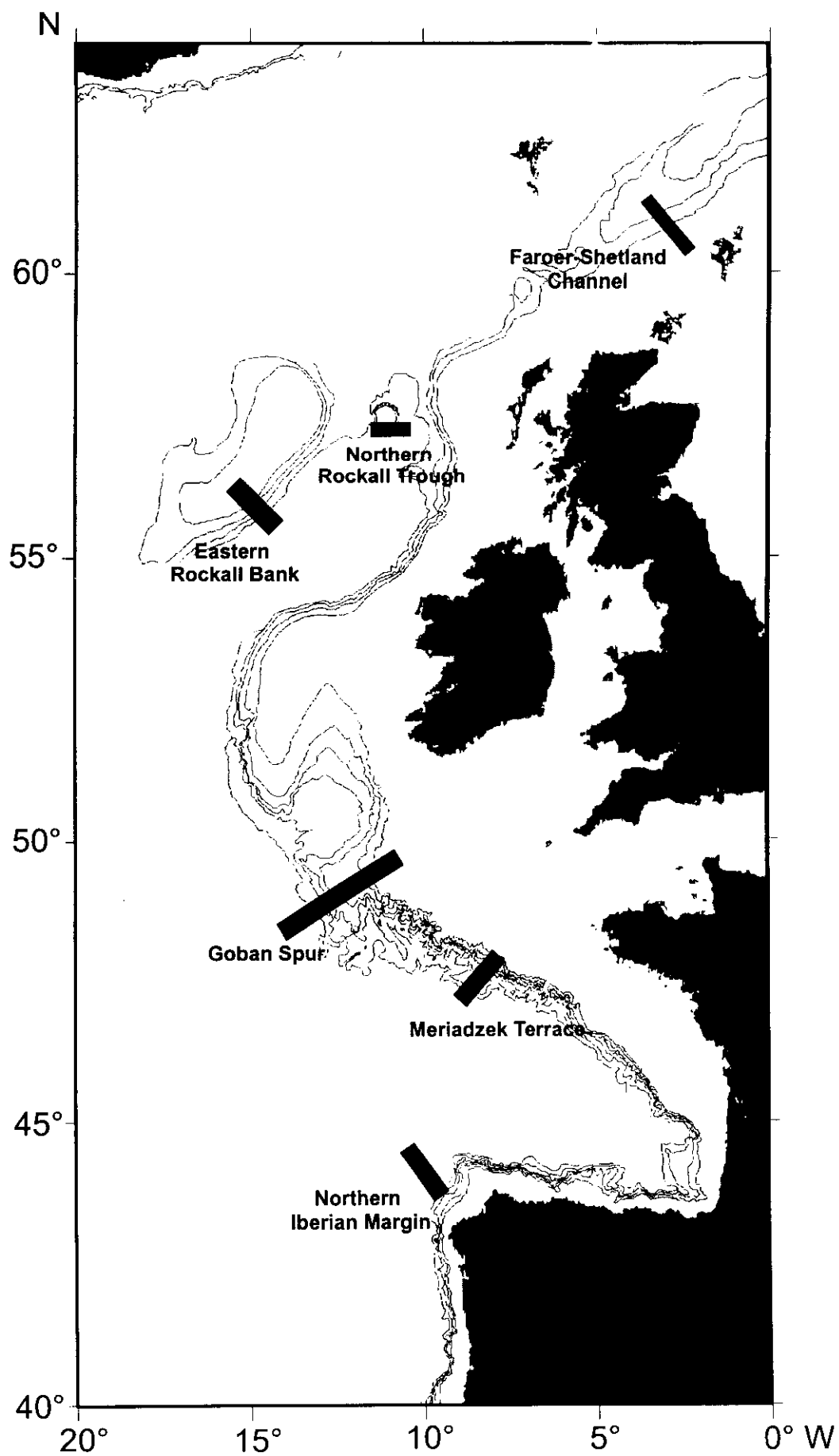
Table 4. Global depth distribution of burial and mineralization rates

Depth range	Area (m)	ω	F_b %	F_c	AEROBIC	DEN	SRR
0-200	7	56	68	52	42	63	64
200-2000	9	30	27	30	30	29	29
> 2000	84	14	5	18	28	8	7

Middelburg et al. Deep Sea Res.



Data points from Fohse et al: Regr. in Oceanogr. (press)
 Regression lines from Middelburg et al.



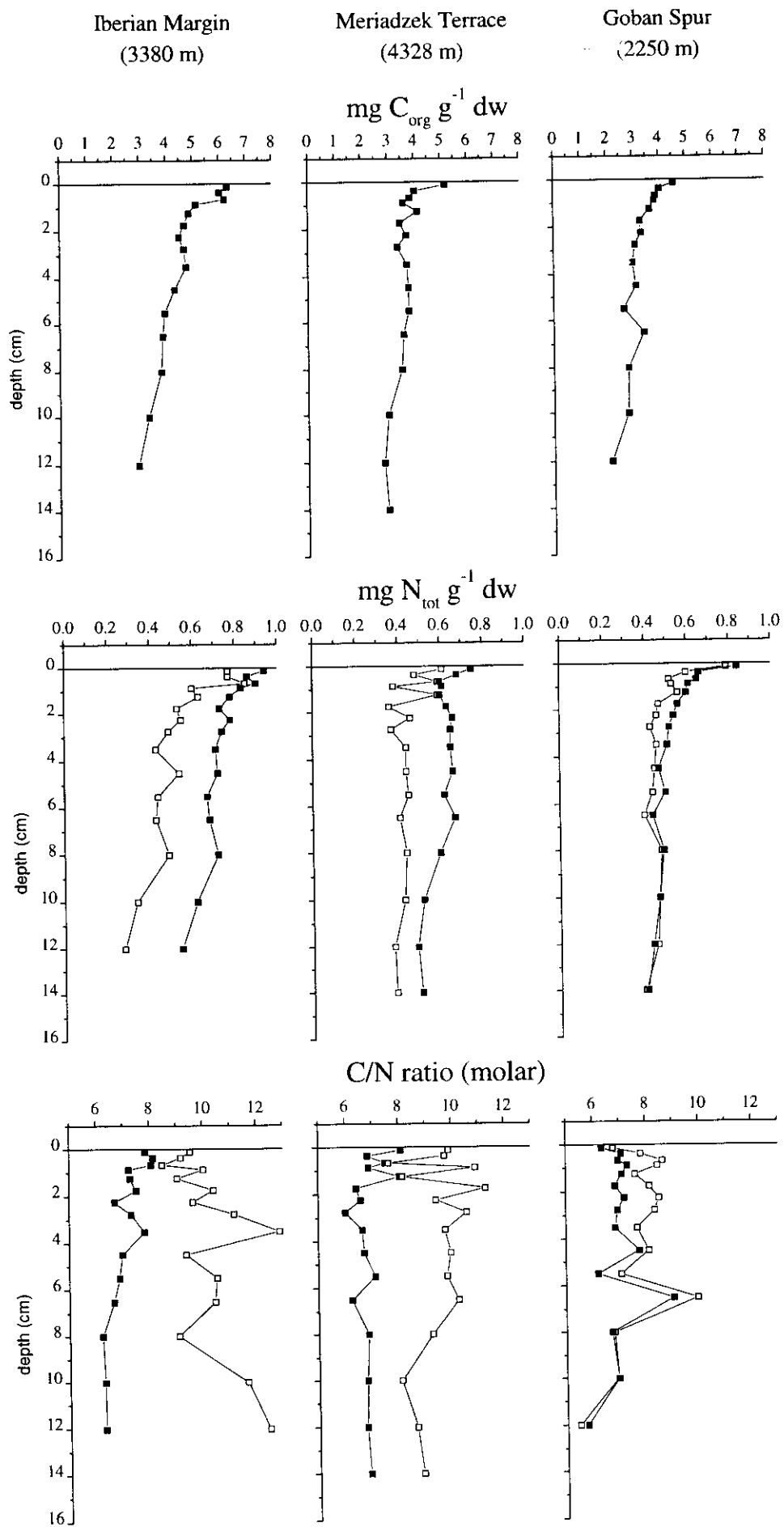


Figure 2
Lohse et al.

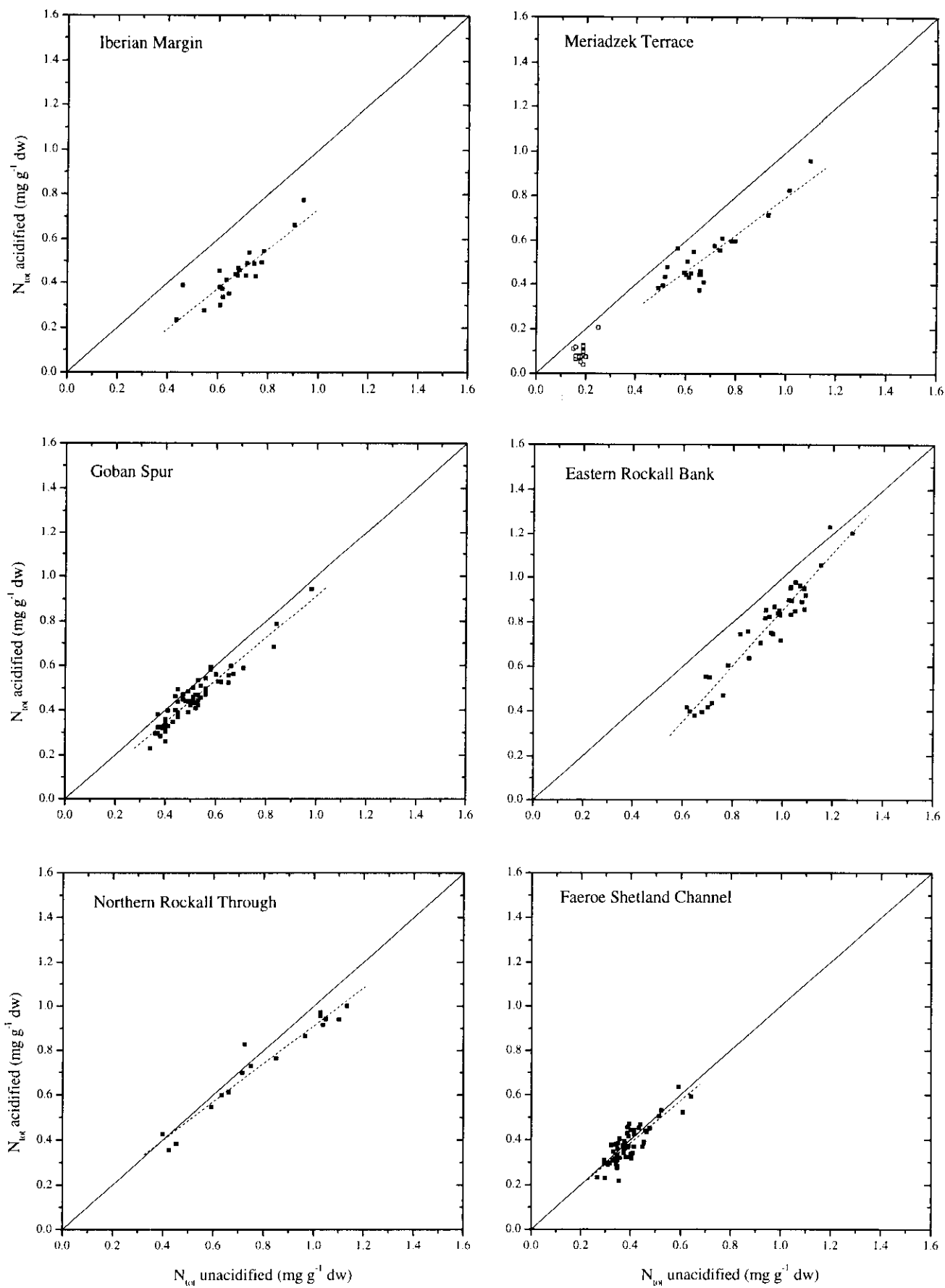


Figure 3 Lohse et al.

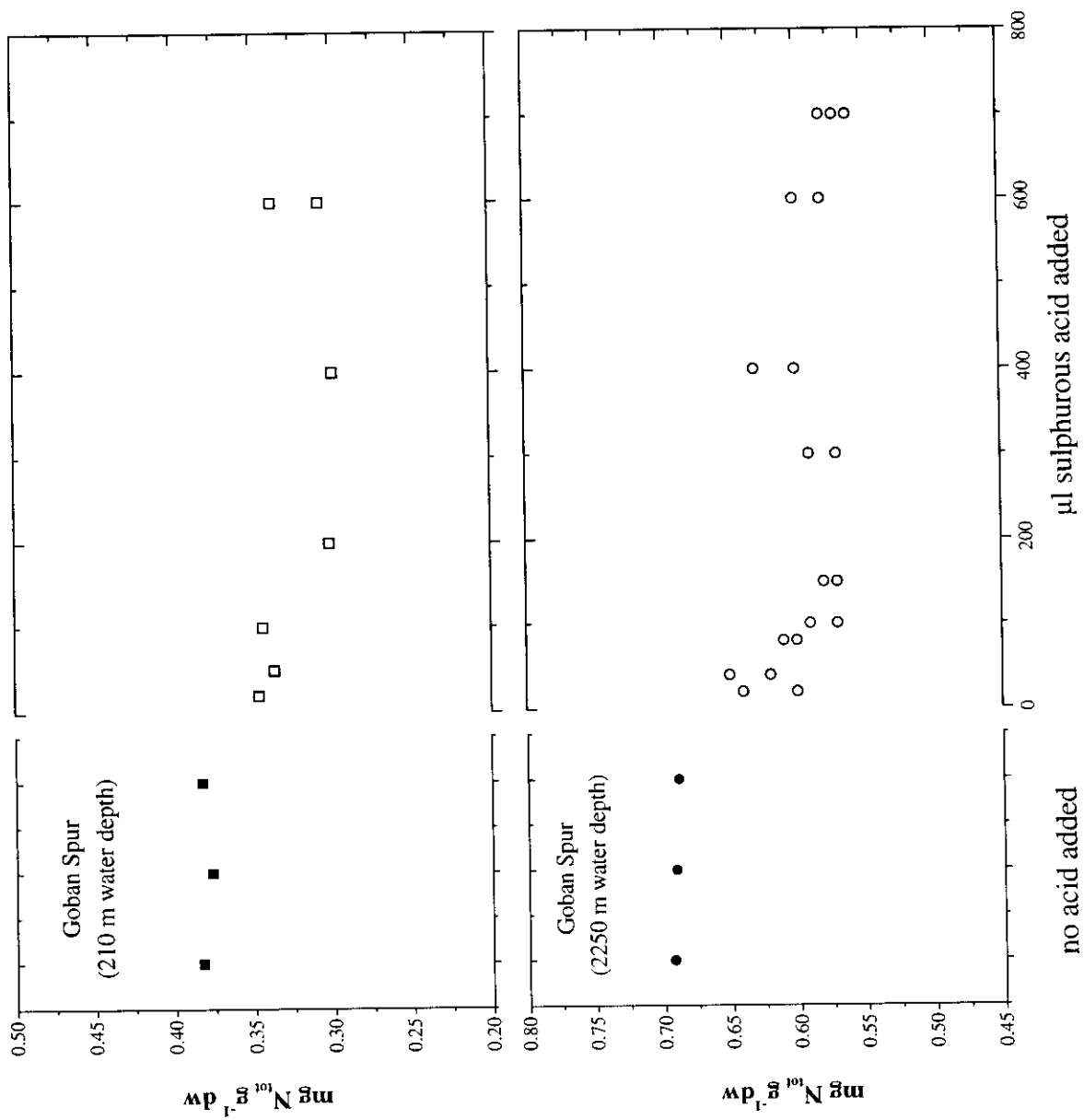


Fig. 4
 Lohse et al.

Water depth	Thickness oxic zone	Sedimentation rate	Exposure time to Oxygen	C _{org} /SFA
(m)	(cm)	(cm x 1000 yr ⁻¹)	(yrs)	(mg / m ²)
600	0.8	2.8	279	0.3
1400	1.4	1.8	841	0.16
1900	9.6	4.3	2242	0.17
3600	14.2	6.0	2367	0.07

