

Organic matter mineralization and nitrogen transformations during stagnation of the Gotland Sea deep water

B. Schneider, IOW

Measurement programme, MAR 2003 – OCT 2007:

Variables:

total CO₂ (C_T,min)

O₂

H₂S

NO₃⁻

NH₄⁺

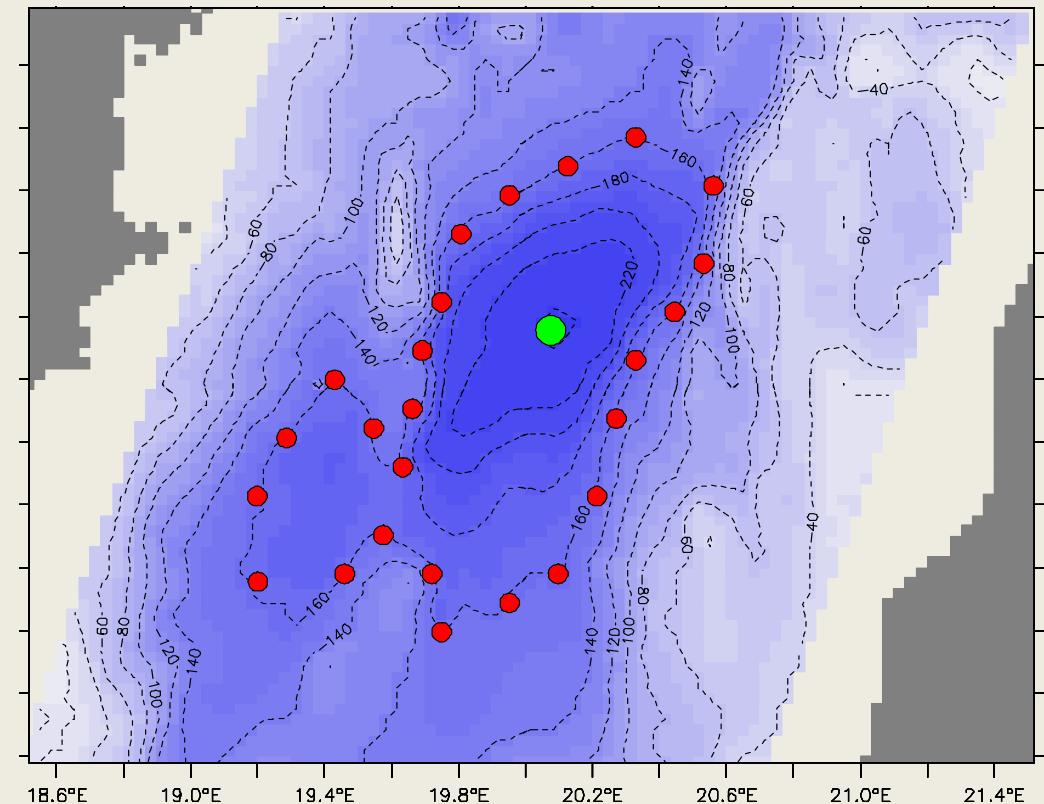
PO₄³⁻

Temporal resolution:

2 – 3 months

Depth resolution:

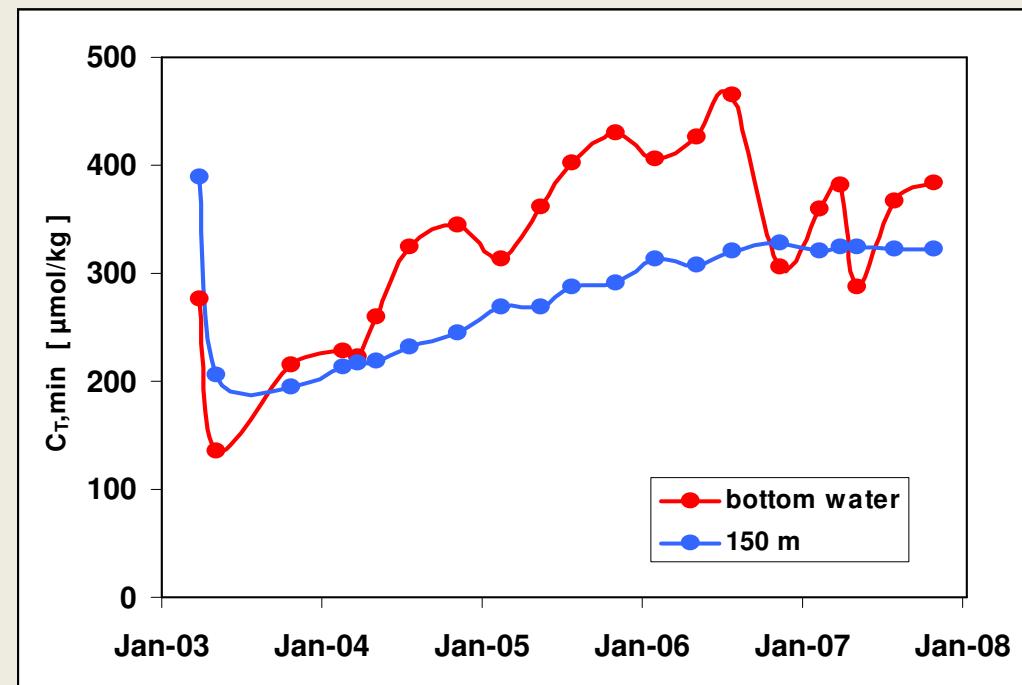
25 m + bottom water



Subdivision in boxes:

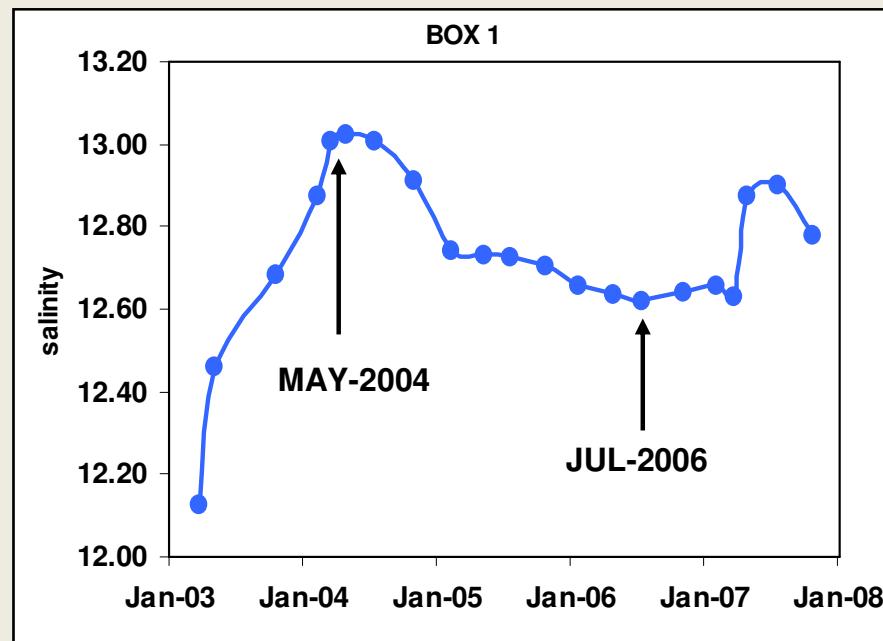
BOX #	Depth [m]	area [km ²]	sediment area [km ²]	volume [km ³]
1	> 225	464	464	3.8
2	200 - 225	1160	696	20.8
3	175 - 200	2120	960	39.7
4	150 - 175	4680	2560	84.9

Time series of the mineralization fraction of total CO₂, C_{T,min}:

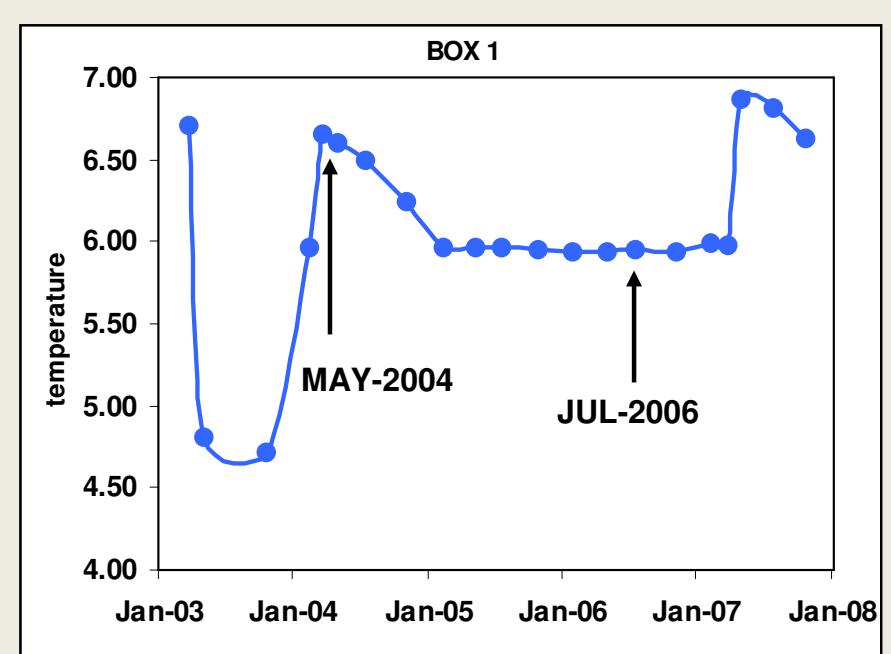


Identification of a period of stagnation:

salinity



temperature



Determination of mixing coefficients, k_{mix}:

$$F_s = k_{\text{mix}} \cdot \frac{\Delta S}{\Delta z}$$

-separately for each time interval;

- gradients: difference in salinity between two neighbouring boxes divided by the distance between the center of the boxes; averaging of the gradients at the beginning and the end of the considered time interval;

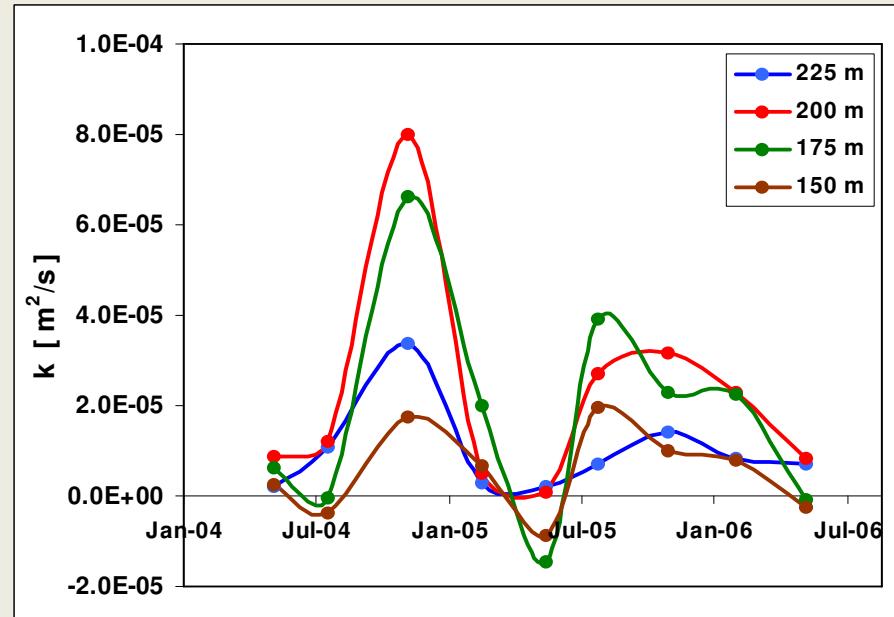
- fluxes: obtained from salt mass balances for each time interval:

$$F_s(1 \rightarrow 2) = (\Delta S_1 \cdot V_1 / A_1) / \Delta t$$

$$F_s(2 \rightarrow 3) = -F(1 \rightarrow 2) + (\Delta S_2 \cdot V_2 / A_2) / \Delta t$$

Mixing coefficients:

- maximum during winter;
- negative values at 175 m and 150 m indicate slight lateral intrusions;



Mass balances for biogeochemical constituents:

$$\frac{\Delta c_n}{\Delta t} = \frac{A_{n-1} \cdot F_{n-1} + A_n \cdot F_n}{V_n} + Q_n$$

n – box number

c – concentration

Δt – time interval

A – area

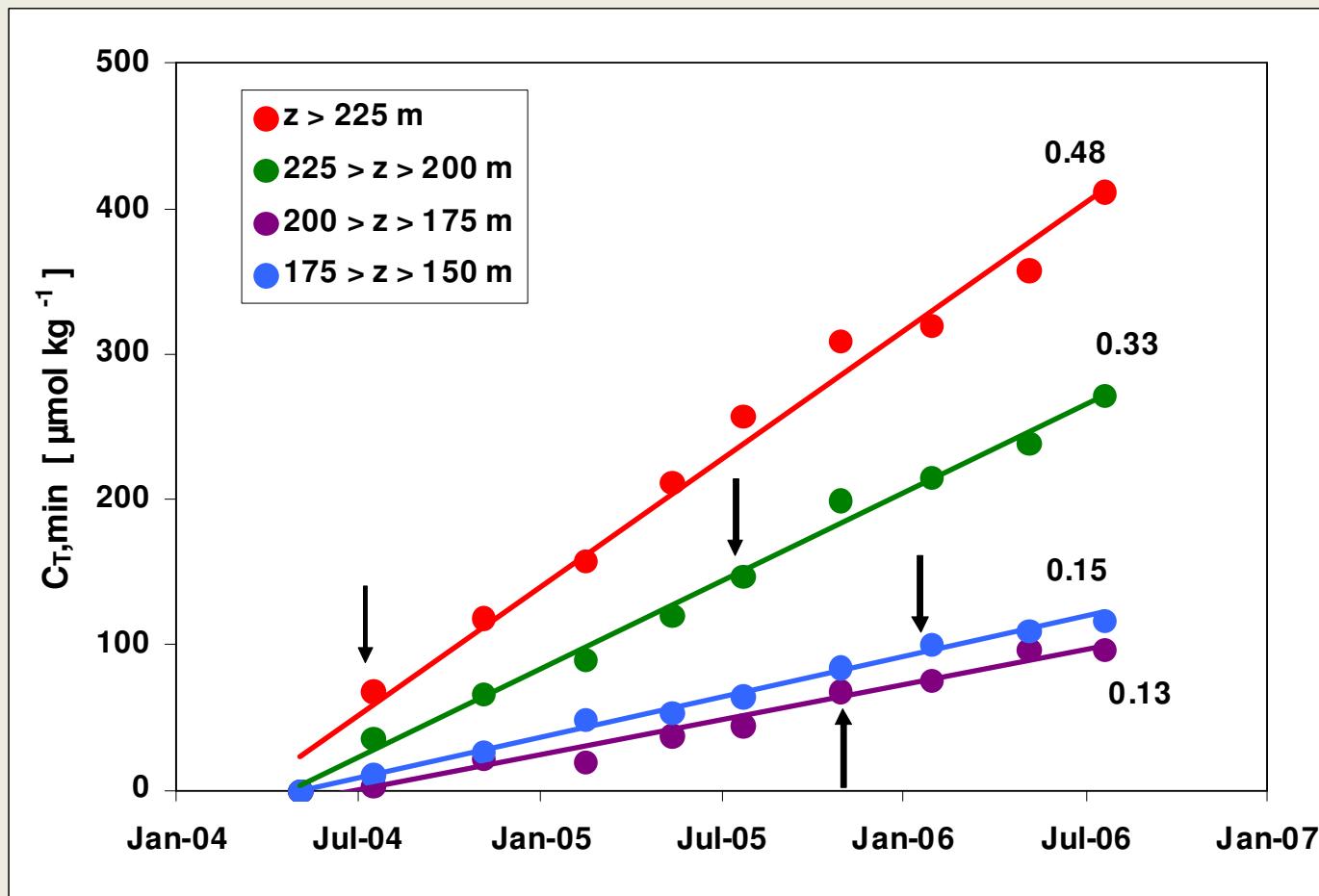
V - volume

F – vertical flux between boxes

Q – sink/source

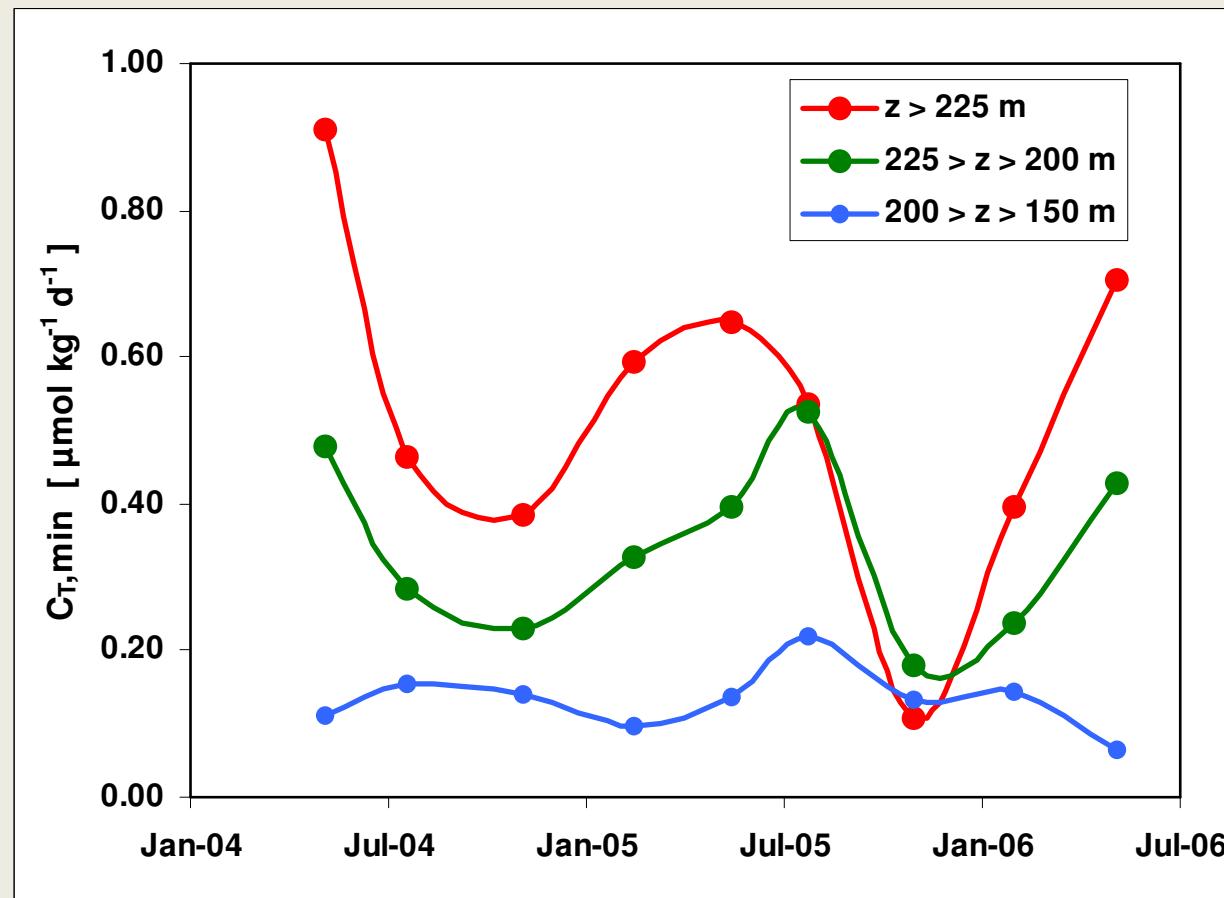
Accumulated C_{T,min} increase during stagnation in different sublayers:

- the arrows indicate the transition to anoxic conditions;
- mean mineralization rates vary between 0.48 and 0.13 ($\mu\text{mol/kg}/\text{d}$);



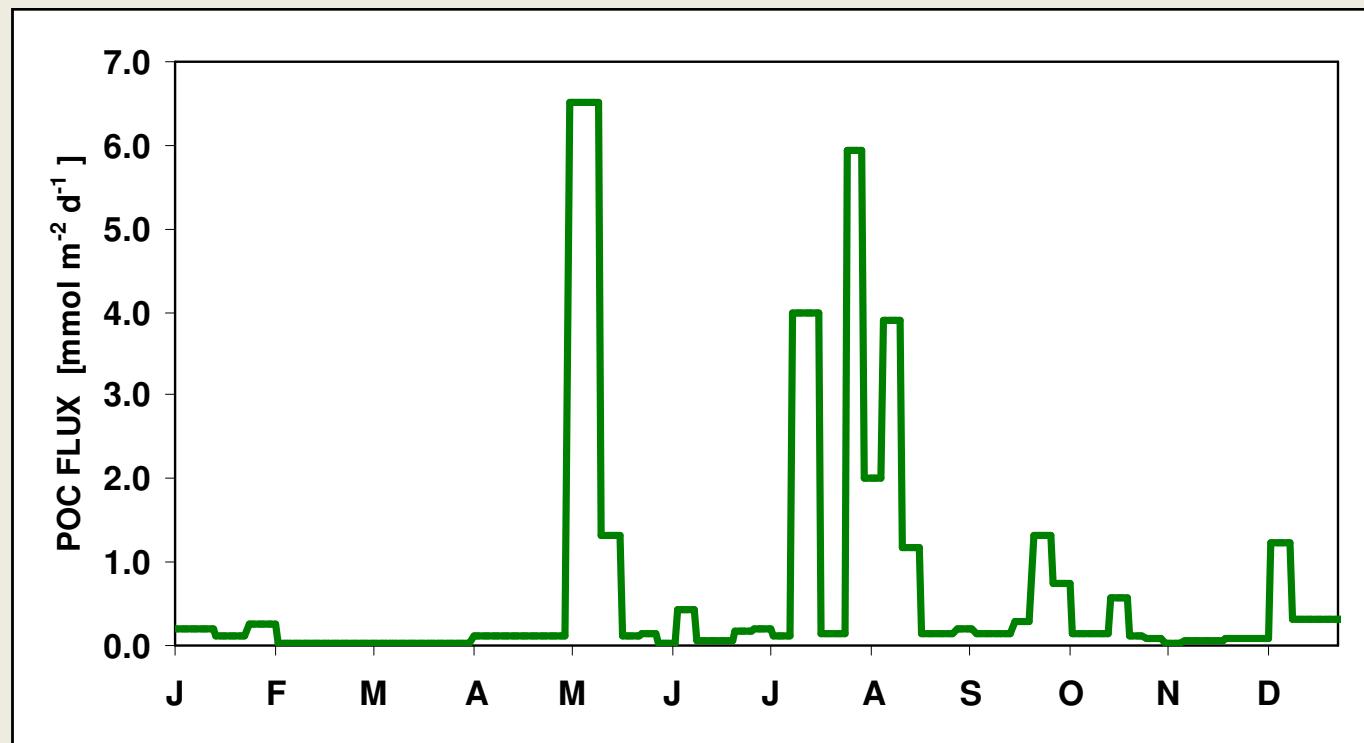
Mineralization rates for individual time intervals:

- maxima during May – July (July – November);
- no seasonal signal above 200 m;



POC fluxes obtained from a sediment trap deployed at 180 m:

- data from 1999;
- data used to estimate the mineralization kinetics;



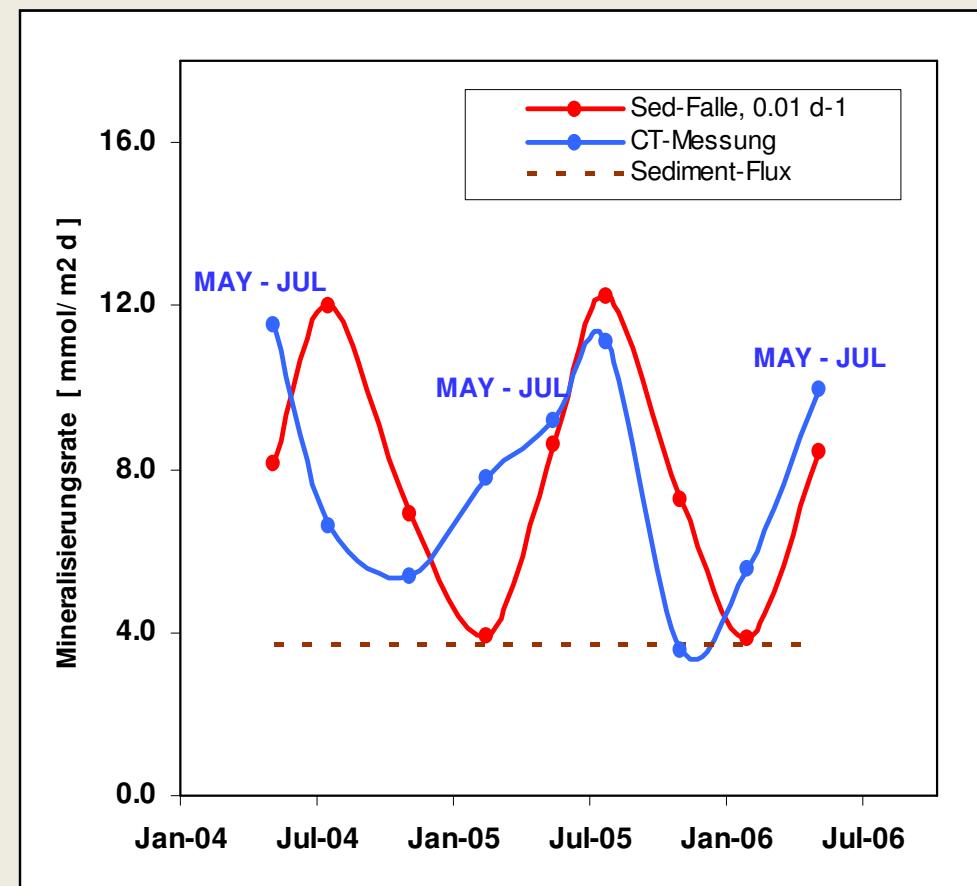
(F. Pollehne, unpublished)

Mineralization kinetics:

$$-\frac{dc_{\text{org}}}{dt} = \frac{dC_T}{dt} = k_{\min} \cdot c_{\text{org}} \cdot c_{\text{oxi}}$$



- modelling the mineralization rates using the POC flux data;
- least square fit yielded best agreement with observed rates for $k_{\min} = 0.01 \text{ d}^{-1}$;
- minima correspond to mineralization of refractory organic matter accumulated in deeper sediment layers;



Redox balance:

		µmol L ⁻¹ y ⁻¹	mmol m ⁻² y ⁻¹	O ₂ equivalents µmol L ⁻¹ y ⁻¹
oxidation	C _{org}	63.6	2024	- 78.9
	NH ₄ ⁺	2.8	89	-2.1
reduction	O ₂	22.3	709	22.3
	SO ₄ ²⁻	20.1	638	40.1
	NO ₃ ⁻	5.1	161	6.3
balance				-12.3

Oxidant demand for the mineralization of organic matter:

- low oxidant consumption in the intial phase of the O₂ depletion/H₂S formation;
- oxidation by MnO₂ ?

