

### Carbon Fractionation and Balance in the Coastal Water of Alexandria Region

A.-R. ABDEL-MOATI, T.-A. ABOUL-KASSIM, N.-M. DOWIDAR and F. EL-NADY

Oceanography Department, Faculty of Science, Alexandria University, Moharam Bay, Alexandria (Egypt)

Carbon is the most important metabolic element in sea water. It occurs in water as a result of precipitation containing increased amounts of  $\text{CO}_2$  produced through fuel combustion,  $\text{CO}_2$  fixation by aquatic plants, from agricultural drainage as organic matter or as major element of waste and sewage discharge. The knowledge of distribution of dissolved organic carbon (DOC) and coexisting particulate organic carbon (POC) is essential for understanding of carbon cycle in sea water. The present work is an attempt to assess the relative importance of land run-off on different carbon species and its contribution to the total carbon budget in a coastal bay off Alexandria falling under sewage stress.

The study area (2.5  $\text{Km}^2$ ) is a semi-circular shallow bay, surrounded by the city, connected to the Mediterranean through two openings. The basin is subjected annually to about  $35 \times 10^6 \text{m}^3$  of unprocessed sewage rendering its flushing rate to be 5 months.

Regular bimonthly sampling during 1985-1986 indicate elevated surface levels of POC reaching 6 mg C/l coinciding with maximum discharge periods as indicated by low salinities. Detrital POC constitutes about 28% of POC. Particulate inorganic carbon (PIC) constituted between 45-49% of total particulate carbon with an average of 4.62 mg/l. On the other hand, the DOC values recorded in the coastal water of Alexandria (average 13.95 mg/l) reflect the highly eutrophic characteristics of water. The organic forms of carbon thus could be used as index for organic pollution derived from sewage discharge.

Generally the dissolved organic fraction exceeds the inorganic by three. The average DOC/POC i.e. 2.4:1 is normal compared with other coastal waters.

The discharged sewage not only affect the carbon in water but raised carbon-sediment levels to 9.11% at areas directly affected by discharge.

The outstanding features of the carbon balance (Figure 1) are:

1. About 325 tons of organic carbon reaches the bay annually through land sources; 70% of which are in particulate form.

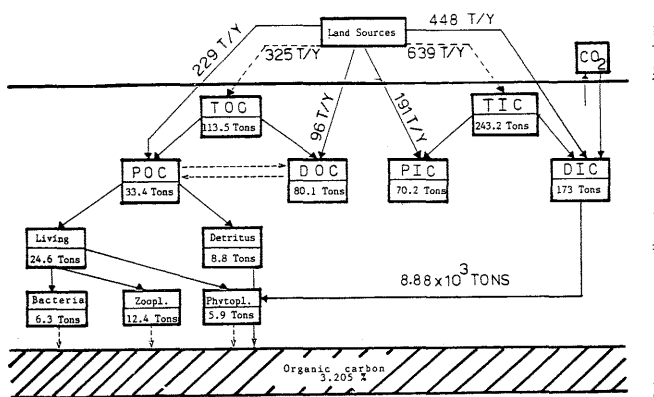


Figure 1. Schematic diagram of carbon input to the Eastern Harbour.

2. The bay receives 639 tons/y of inorganic carbon of which the particulate fraction forms only 30%.
3. About 15% of POC influx existed in the water column as living (74%) while about 37% of PIC influx is retained in suspension.
4. 85% of the inflowing DOC exists in the water column. The equilibrium shift of  $\text{POC} \rightleftharpoons \text{DOC}$  towards DOC may substitute the loss in DOC during oxidation processes.
5. About  $8.88 \times 10^3$  tons carbon are fixed annually by phytoplankton. Atmospheric  $\text{CO}_2$  input  $4.4 \times 10^3$  tons/y could substitute a significant part of this uptake rate.
6. The short residence time of the bay water (5 months) leads to a mismatch between the inflowing carbon and that actually present in the bay as well as a considerable differences in the proportionality of different carbon species.

### Land Run-Off as a Source of Nitrogen in the Marine Coastal Environment of Alexandria

F.-E. EL-NADY, T.-A. ABOUL-KASSIM, N.-M. DOWIDAR and A.-R. ABDEL-MOATI

Oceanography Department, Faculty of Science, Alexandria University, Moharam Bay, Alexandria (Egypt)

**INTRODUCTION:** Nitrogen is one of the biologically important elements in the aquatic habitats. In addition to dissolved molecular nitrogen, sea water contains low, but extremely important, concentrations of inorganic and organic nitrogen. The present work deals with the concentrations of the different nitrogen species as well as their contribution to the total nitrogen budget in one of the most polluted basins off Alexandria, the Eastern Harbour (E.H.).

**MATERIAL AND METHODS:** During the period 1985-1986, sampling was carried out at regular bimonthly intervals in a semi-closed bay, connected to the Mediterranean through two openings. The basin is subjected to about  $35 \times 10^6 \text{m}^3$  of unprocessed sewage, rendering its flushing rate to be 5 months. Nitrate, nitrite and ammonia were determined according to Strickland and Parsons (1972). Total dissolved nitrogen (TDN) and total nitrogen (TN) were determined on filtered and unfiltered samples using the technique described by Koroleff (1977) and modified by Valderrama (1981). Dissolved organic nitrogen (DON) and particulate nitrogen (PN) were estimated by calculations. The uptake rate of  $\text{NO}_3^-$  &  $\text{NH}_3$  by phytoplankton of the harbour water was determined using the procedure of Eppley et al. (1969). The flux of nitrogen from the harbour sediments was measured following the method of Hargrave and Connolly (1978). Organic nitrogen in sediments was determined according to Niederl and Niederl (1942).

**RESULTS AND DISCUSSION:** Nitrate is the final oxidation product of nitrogen compounds in sea water. During the study period, its concentration in the E.H. was comparatively high, the annual averages being  $6.791 \pm 4.654$  and  $4.826 \pm 2.964 \mu\text{g at/l}$  for both surface and bottom waters, respectively. The nitrite concentrations were much lower than that of nitrate (The averages being 0.949 and  $0.746 \mu\text{g at/l}$  for both surface & bottom waters, respectively). The importance of waste water discharged into the harbour as a source of ammonia was found from the inverse correlation between ammonia and salinity ( $P < 0.001$ ). In spite of shallowness of the E.H., ammonia concentration was relatively high, varying between  $0.975 - 11.456 \mu\text{g at/l}$  (at the surface) and  $0.480 - 12.334 \mu\text{g at/l}$  near the bottom. The observed correlation ( $P < 0.001$ ) between  $\text{NO}_3^-$  &  $\text{NO}_2^-$  content and its insignificance with ammonia indicated that nitrate reduction rather than ammonia oxidation is a major source of nitrite.

Dissolved organic nitrogen (DON) was comparatively higher (annual average  $11.866 \pm 6.129 \mu\text{g at/l}$ ) than that of DIN (average  $10.06 \pm 4.864 \mu\text{g at/l}$ ). This is probably due to being assimilated by aquatic organisms at a much lower rate than inorganic forms or being resistant to bacterial attack, remaining in the water or sinking to the bottom (Riley and Chester, 1971). The correlations between DON with  $\text{Chl a}$  ( $P < 0.001$ ) and living biomass represented by ATP ( $P < 0.001$ ), confirmed the important role of living organisms as a source of DON.

High concentrations of PN occurred in summer, coinciding with the periods of maximum sewage discharge and chlorophyll *a* biomass (Aboul-

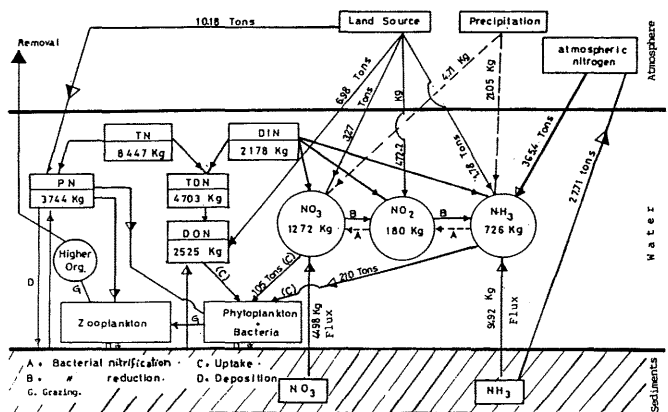


Figure 1: The Nitrogen Balance in the Eastern Harbour of Alexandria.

Kassim, 1987). The highly significant correlations between PN with POM ( $P < 0.001$ ) and salinity ( $P < 0.001$ ) are evidences for the role of phytoplankton and sewage discharge as important sources of PN.

On the average, most of the nitrogen budget of the E.H. (59%) was in the dissolved form. DON was more abundant, constituting about 54% of TDN. Nitrate was the most abundant form of DIN forming 55.7%, followed by ammonia 37.1% and nitrite 7.2%. The mean atomic ratios of the different nitrogen and phosphorus forms, i.e.  $\text{NO}_3^-/\text{DIP}$ ,  $\text{DIN}/\text{DIP}$ ,  $\text{DON}/\text{DOP}$ ,  $\text{TDN}/\text{TDP}$ ,  $\text{PN}/\text{PP}$  and  $\text{TN}/\text{TP}$  were as follows: 13.62:1, 23.97:1, 24.52:1, 22.67:1, 11.33:1 and 15.85:1, respectively.

Within the values given in figure 1, the total input of all nitrogen compounds from land sources, flux from sediments, precipitation and nitrogen fixation amounted to 402 Tons/year. This amount exceeds that due to uptake, denitrification from bottom sediments and the amount present in the harbour environment by 51 Tons/year.

#### REFERENCES:

- Aboul-Kassim, T.A. (1987). M.Sc. Thesis, Fac. of Sci., Alex. Univ., Egypt.  
 Eppley, R.W., J.N. Rogers & J. McCarthy (1969). *Limnol. Oceanogr.* 14: 912-20.  
 Hargrave, B.T. and C.F. Connolly (1978). *Limnol. Oceanogr.* 23: 1005-1010.  
 Koroleff, F. (1977). In: Grasshoff, K. (ed.). *Annex. Interim Commission of the Protection of the Baltic Sea.*  
 Niederl, J.B. and V. Niederl (1942). *Micromethods of quantitative organic analysis* (2nd ed.), John Wiley & Sons, New York, 374 pp.  
 Riley, J.P. and R. Chester (1971). *Marine chemistry*. Academic Press, 465 pp.  
 Strickland, J.D.H. and T.R. Parsons (1972). *Fish. Res. Bd. Canada, Bull.* 167, 2nd ed., 310 pp.  
 Valderrama, J.C. (1981). *Marine Chemistry*, 10: 109-122.