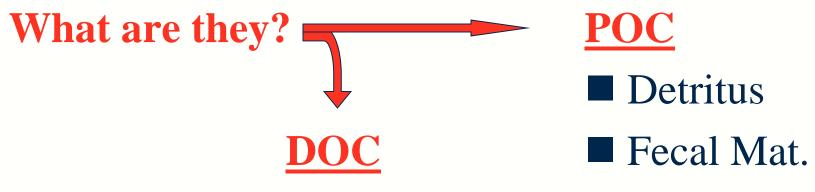
Chemical Oceanography Organic Materials II

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Organic Materials in Marine Environment



- Biological molecules (lipids, proteins, carbohydrates, etc.)
- Hydrocarbons
- Humic Materials (=other stuff)

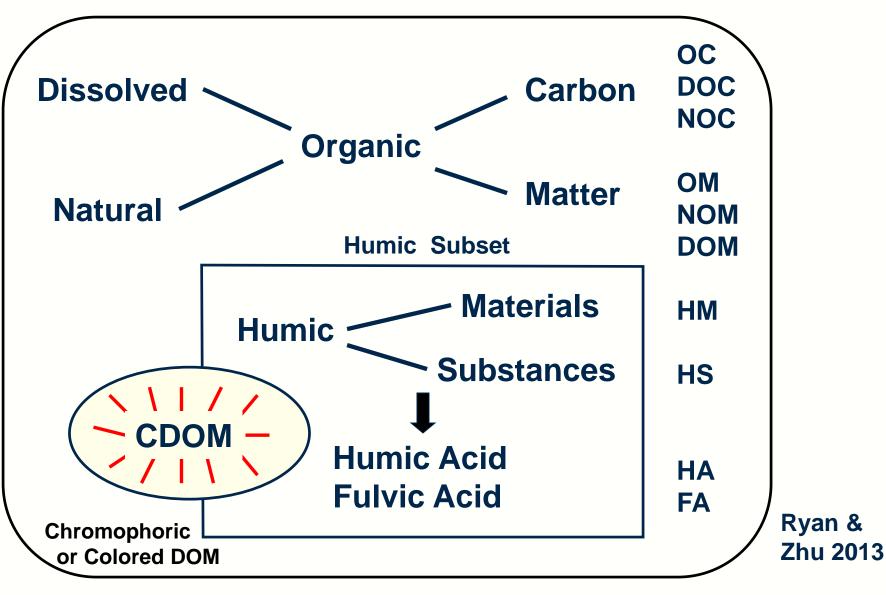
| Average Concentrations of O in Baltic and North S | | |
|--|--|---------------------|
| Components | Concentration $(\mu g C liter^{-1})$ | |
| Free amino acids | 10 | |
| Combined amino acids | 50 (to 100?) | |
| Free sugars | 20 | |
| Combined sugars | 200 | |
| Fatty acids | 10 | |
| Phenols | 2 | |
| Sterols | 0.2 | |
| Vitamins | 0.006 | What is this stuff? |
| Ketones | 10 | |
| Aldehydes | 5 | |
| Hydrocarbons | 5 | |
| Urea | 10 | Morel, 1983 |
| Uronic acids | 18 | |
| Approximate identified total | 340 μ g C liter ⁻¹ | |
| Approximate total | $\begin{array}{c} 340 \ \mu \text{g C liter}^{-1} \\ 4000 \ \mu \text{g C liter}^{-1} \end{array}$ | 3 |
| | | 5 |

| Table 8.4. Molecular-level methods for different types of organic substance | | | | | | |
|---|--------------|-------|--------------------------|---------------------|-------------------------|-----------------------|
| Method | Class | Types | ^a Preparation | ^b Chrom. | ^c Derivative | ^d Detector |
| Hydrocarbons | Lipid | >100 | NPSE | GC | None | FI |
| Fatty acids | Lipid | > 100 | Basic Hy | GC | ME/TMS | FI |
| Fatty alcohols | Lipid | с.30 | Basic Hy | GC | ME/TMS | FI |
| Sterols | Lipid | >100 | Basic Hy | GC | ME/TMS | FI |
| Alkenones | Lipid | с.10 | NPSE | GC | None | FI |
| Chlorophylls | Pigment | с.20 | NPSE | LC | None | Flu |
| Carotenoids | Pigment | с.50 | NPSE | LC | None | UV |
| Amino acids | Amine | с.20 | Acid Hy | LC/GC | OPA | Flu/Fl |
| Nucleic acids | Nucleotide | 4 | Isolation | LC | None/OPA | UV |
| Neutral sugars | Carbohydrate | 20 | Acid Hy | IC/GC | None/TMS | PA/FI |
| Acidic sugars | Carbohydrate | c.10 | Acid Hy | IC/GC | None/TMS | PA/FI |
| Lignin phenols | Phenol | с.30 | CuO-NaOH | GC/LC | None/TMS | FI |
| Tannins | Phenol | с.20 | Acid Hy | GC | PHL/TMS | FI |
| Cutin acids | Polyester | с.20 | MeOH- | GC | ME/TMS | FI |
| Pyrolysis-GC/MS | General | > 100 | Pyrolysis | GC | None | MS |
| TMAH Chemolysis | General | > 100 | TMAH-Heat | GC | ME | FI |
| CuO/NaOH | General | >100 | CuO-NaOH | GC | TMS | FI |

Emerson & Hedges 2010

Flame Ionization (FI), Fluorescence (Flu), Pulsed Amperometric (PA) 4

All Dissolved Organic Compounds



| 10 | 20 | 30 | 40 | 50 | 60 | 70 | 80 | 90 | 100 % |
|----|----|-----------------|---------------------------------------|-----------|--|---|---|---|--|
| 1 | 1 | . 4 | 1 | 122 | 1 | 1 | | 4 | |
| | | | F | ulvic aci | d | | | | |
| | | | Н | umic aci | id | | Carlogu | | 1 |
| | | | H | lydrophi | lic acid | | | | |
| | Ca | rbohydr | ates | | | | | | |
|] | Ca | rboxylic | acids | S | imple | | | | |
| | Am | iino acio | ls | соп | npounds | | | | |
| | Hy | drocarb | ons | | | | | | |
| | | Ca Ca Arr | Carbohydr Carboxylic Amino acid | F | Fulvic aci Fulvic aci Humic ac Hydrophi Carbohydrates Carboxylic acids Scorr | Fulvic acid Humic acid Hydrophilic acid Carbohydrates Carboxylic acids Amino acids | Fulvic acid H Bubs Humic acid Hydrophilic acid Carbohydrates Carboxylic acids Amino acids | Fulvic acid Humic substances Humic acid Hydrophilic acid Carbohydrates Carboxylic acids Amino acids | Fulvic acid Humic substances Humic acid Humic acid Hydrophilic acid Hydrophilic acid Carbohydrates Simple compounds Amino acids Simple compounds |

Libes, 1992

20

FIGURE 23.13. Composition of dissolved organic carbon in average river water with a DOC concentration of 5 mg/L. Source: From Organic Geochemistry of Natural Waters, E. M. Thurman, copyright © 1985 by Kluwer Academic Publishers, Dordrecht, The Netherlands. Reprinted by permission.

Humic Materials

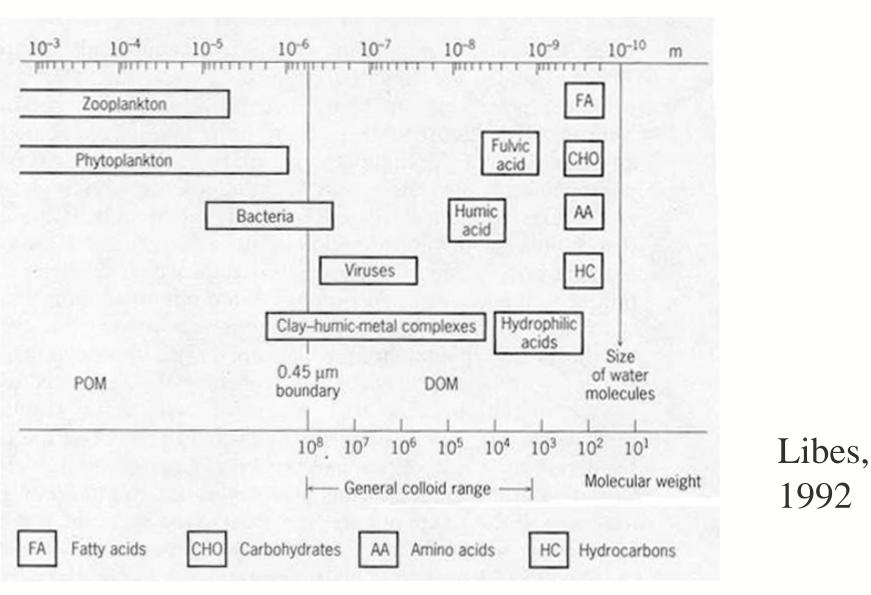
- **H** Complex natural organic molecules
- **#** Properties & importance understood
- **#** Some structural components known
- Exact chemical nature or exact structure unknown because:
 - Complexity
 - Heterogeneity
 - Concentrations

Deficiencies in analytical

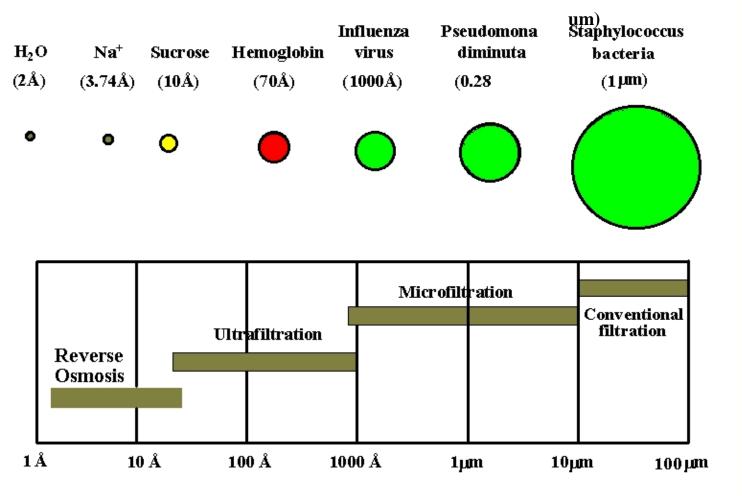
techniques

Interfering species

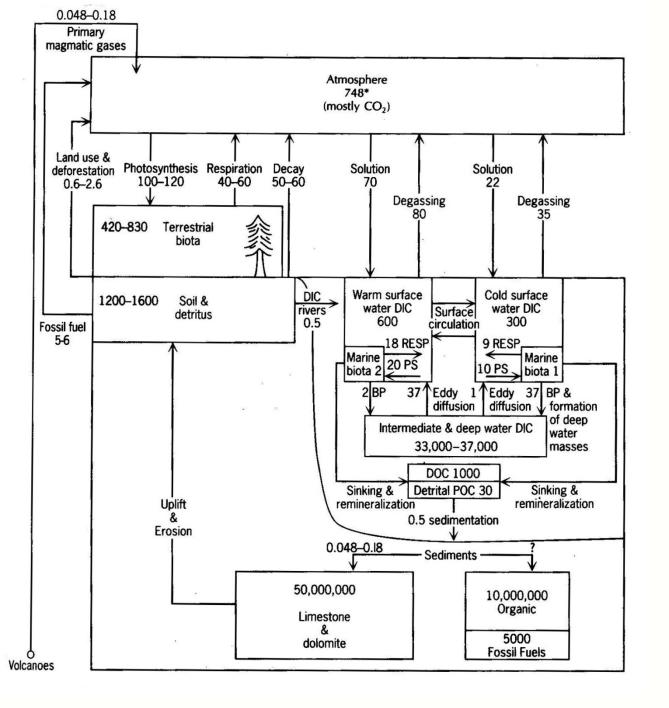
Organic Carbon Continuum



PORE SIZE OF FILTRATION PROCESSES



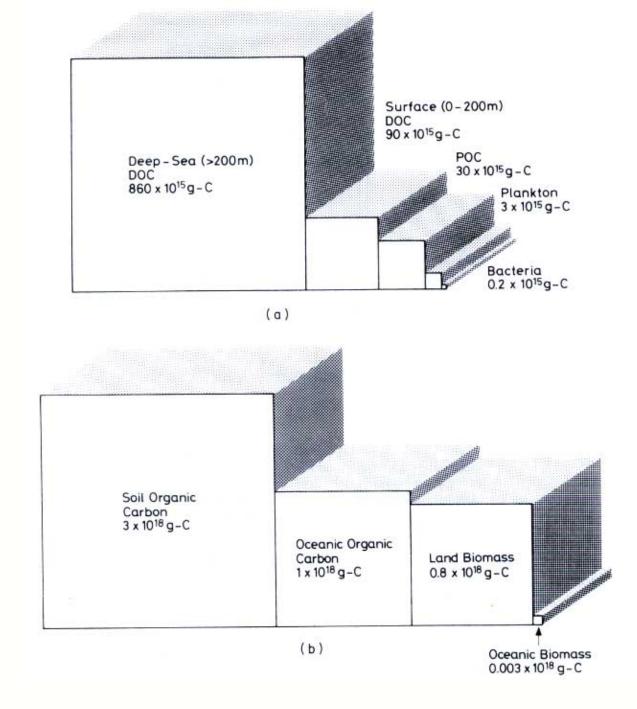
Pore Diameter



Carbon Cycle Libes, 1992

Inventories in 10^{15} g C = BMT

Fluxes (arrows) 10¹⁵ g C/yr



Distribution of Organic Carbon

- (a) Major compartments in the global ocean
- (b) Major compartments for the planet

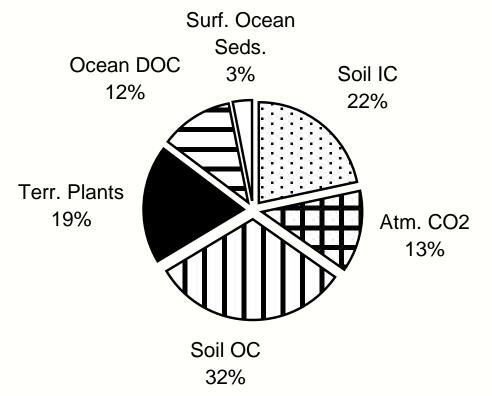
Cauwet, 1978

Major reservoirs of organic and inorganic carbon

| D | Amount |
|--------------------------------|------------------------|
| Reservoir type | (10 ¹⁸ g C) |
| Sedimentary Rocks | |
| Inorganic (Carbonates) | 60,000 |
| • Organic (e.g. kerogen, coal) | 15,000 |
| | |
| Active (surficial) pools | |
| Inorganic | |
| • Marine DIC | 38 |
| Soil Carbonate | 1.1 |
| • Atmospheric CO ₂ | 0.66 |
| Organic | |
| • Soil humus * | 1.6 |
| Land plant tissues | 0.95 |
| Seawater DOC | 0.60 |
| Surface marine sediments | 0.15 |

After Hedges, 1992; * pre-anthropogenic values.

Active Carbon Reservoirs (excluding Ocean DIC)



Organic Compounds in Marine Environment

- **Where do they come from?**
- **What are they?**
- Bio & Geo
 Hydrocarbons
 - Carbohydrates (polysaccharides), sugars
 - Lipids, fats, waxes, oils, fatty acids
 - Pigments

Bio

?

- Nucleic acids, RNA, DNA
- Amino acids, polypeptides, proteins, enzymes
- Low molecular weight carboxylic acids
- Humic Substances

| Table 8.1. Number of structural isomers for alkanes of increasing carbon number | | | | | |
|---|-------------------|---------------------------------|-------------------|--|--|
| Formula | Number of isomers | Formula | Number of isomers | | |
| $C_{6}H_{14}$ | 5 | $C_{10}H_{22}$ | 75 | | |
| C_7H_{16} | 9 | $C_{15}H_{32}$ | 4347 | | |
| C_8H_{18} | 18 | $C_{20}H_{42}$ | 366 319 | | |
| C ₉ H ₂₀ | 35 | C ₃₀ H ₆₂ | 4 846 763 | | |

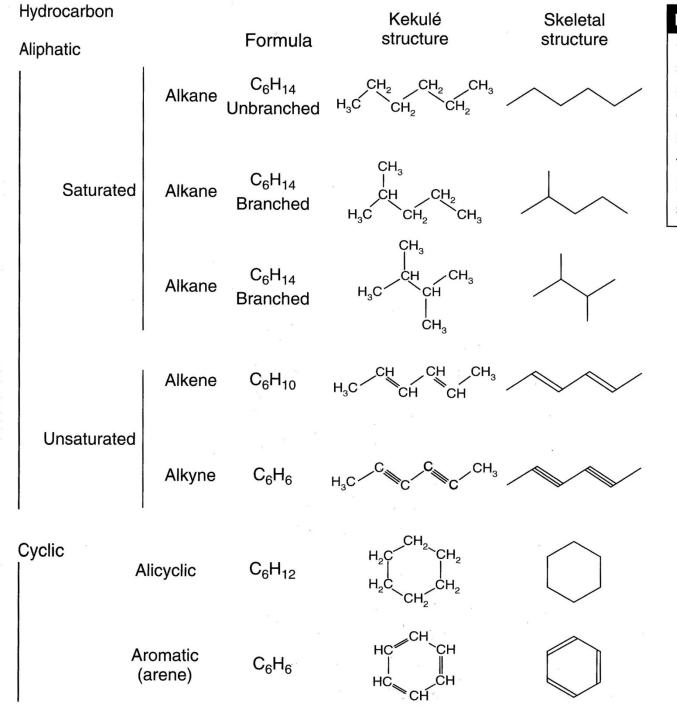


Figure 8.1. Hydrocarbons are classified by whether their carbon structure includes rings, multiple bonds, or branches. Classification of hydrocarbon structural families is shown here with the chemical formula. Structures are shown as Kekulé (also called line-bond) and skeletal structures.

Organic Compounds in Marine Environment

- **Where do they come from?**
- **What are they?**
- Bio & Geo
 Hydrocarbons
 - Carbohydrates (polysaccharides), sugars
 - Lipids, fats, waxes, oils, fatty acids
 - Pigments

Bio

?

- Nucleic acids, RNA, DNA
- Amino acids, polypeptides, proteins, enzymes
- Low molecular weight carboxylic acids
- Humic Substances

Organic Carbon Inputs to the Ocean

Allochthonous = formed externally (*ex situ*) **#** Autochthonous = formed internally (*in situ*) Most Marine Humic Material is formed *in situ* through both biotic & abiotic processes Some Humic Material (i.e., coastal) is introduced from terrestrial sources (formed on land)

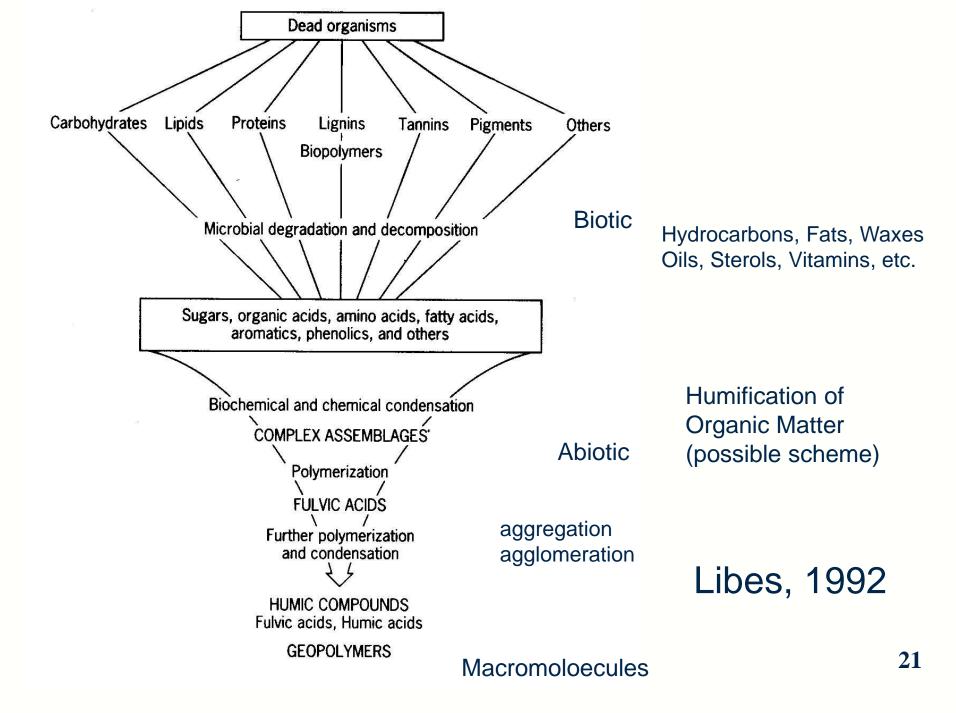
Transformation of DOC

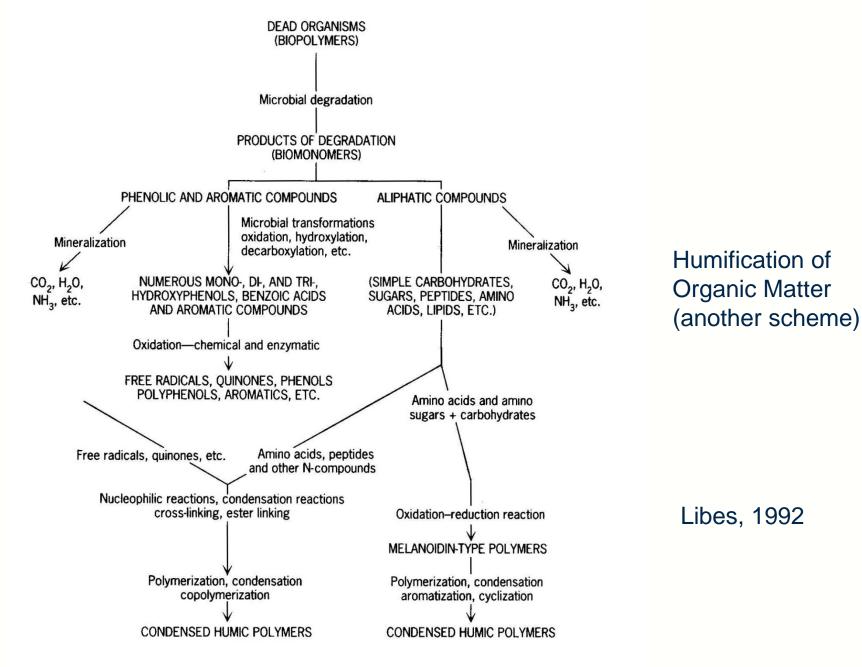
Biological molecules are
labile = readily broken down
or degraded quickly

- By-products of this breakdown (substances not completely remineralized) can react with other organic compounds in a process called Humification or Early Diagenesis
- **#** This results in non-labile **Humic Materials**
- Humics may degrade slowly or be removed to the sediments (refractory or non-labile)

Transformation of DOC

- These processes occur inthe water column, in sediments, & in soils
- **#** Humification is the first step, fast, aerobic
- Fossilization or carbonification occur more slowly on geologic time scales, anaerobicly, after burial in sediments
- **#** Diagenesis, Catagenesis, Metagenesis



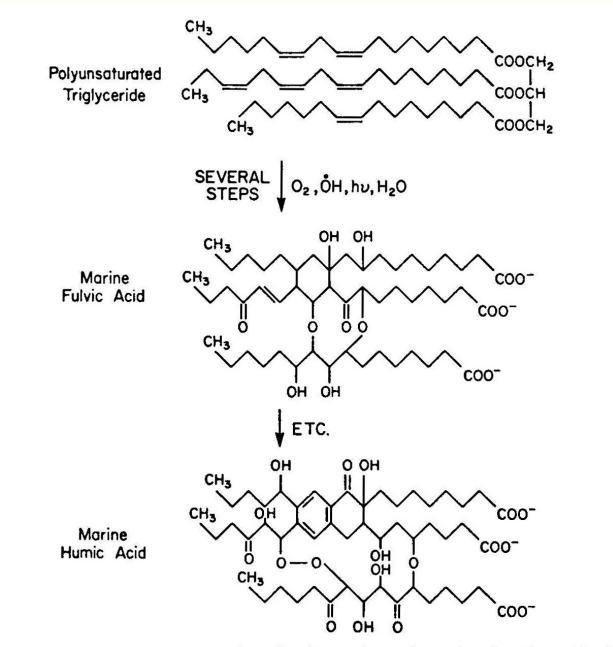


(GEOPOLYMERS)

| Group | Structure | pK _a | Hydrolysis Products | Exchange H ? |
|----------|--|-----------------|------------------------|-----------------|
| Alcohol | -С- О-Н | 12 | None | Yes |
| Phenol | О-о-н | 10 | None | Yes |
| Ether | $-\mathbf{c} - \mathbf{c} -$ | | None | |
| Aldehyde | о | | None | No |
| Ketone | -c - c - c - c - c - c - c - c - c - c | | None | |
| Carboxyl | о С-С-О-Н | 5 | None | Yes |
| Ester | $-\stackrel{\mathbf{O}}{\overset{\mathbf{O}}}{\overset{\mathbf{O}}{\overset{\mathcal{O}}{$ | | Carboxyl + Alcohol | |
| Amine | $-\mathbf{\dot{C}}-\mathbf{N}$ | 10 | None | Yes |
| Amide | 0 −C−C−N | | Carboxyl + Amine | Yes |

Emerson & Hedges Figure 8.2

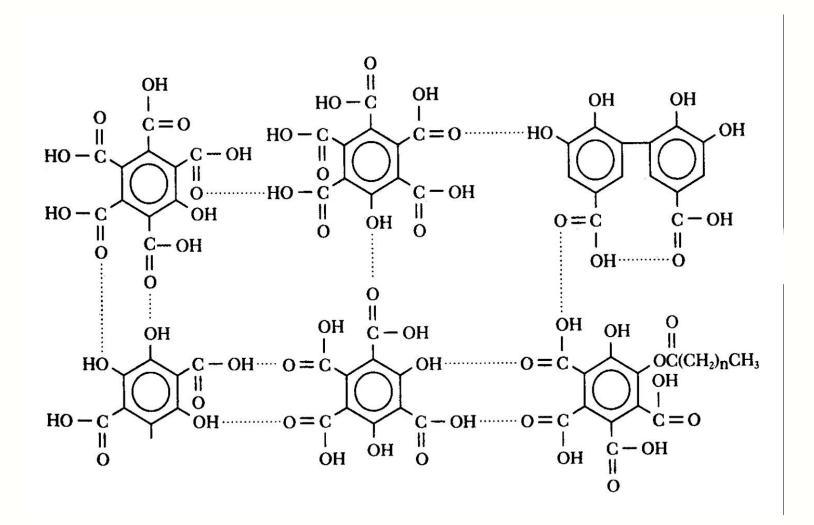
23



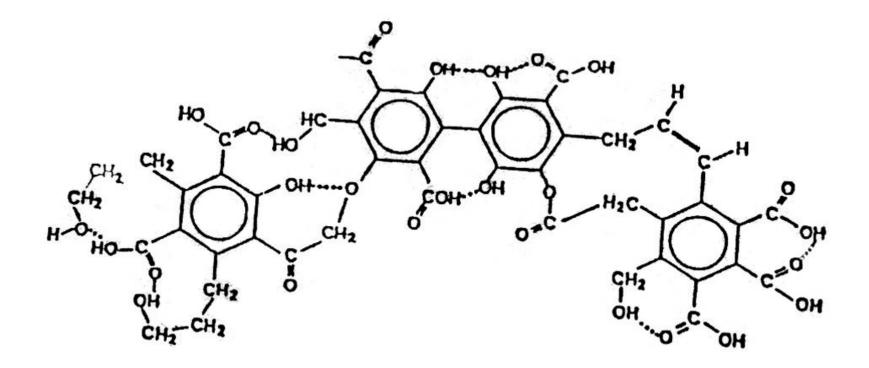
Morel & Hering, 1993 See also Emerson & Hedges Figure 8.11 & 8.12

Figure 6.13 A possible pathway for the formation of marine humic acids from a triglyceride. From Harvey et al., 1983.

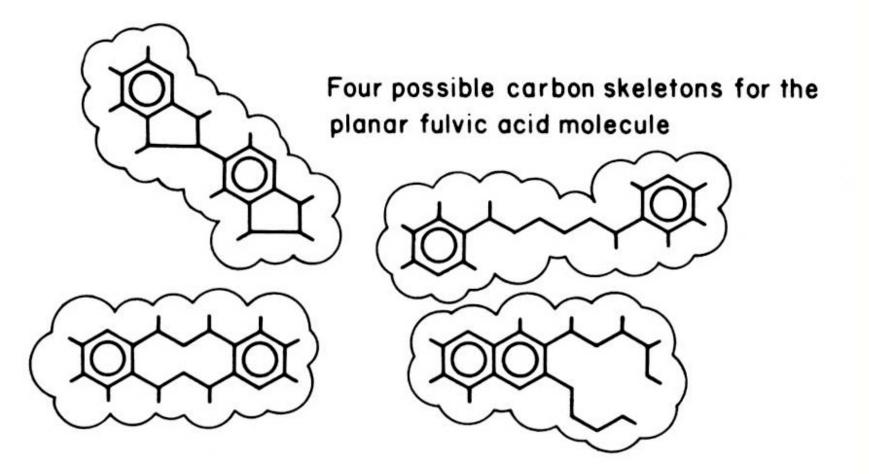
24



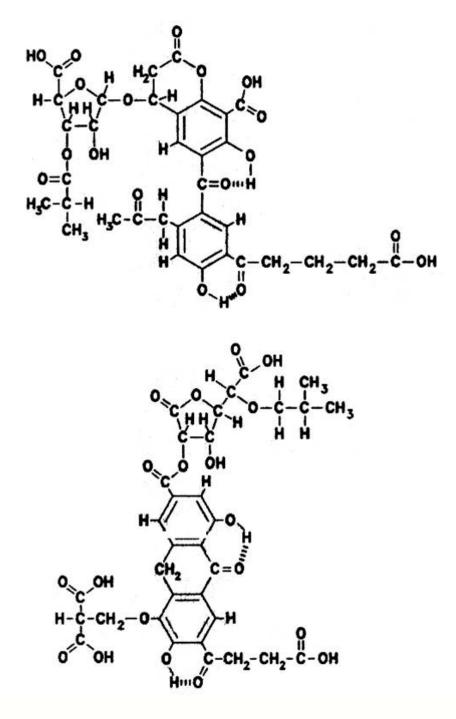
Humic Structure Proposed by Schnitzer (Rashid 1985) 25



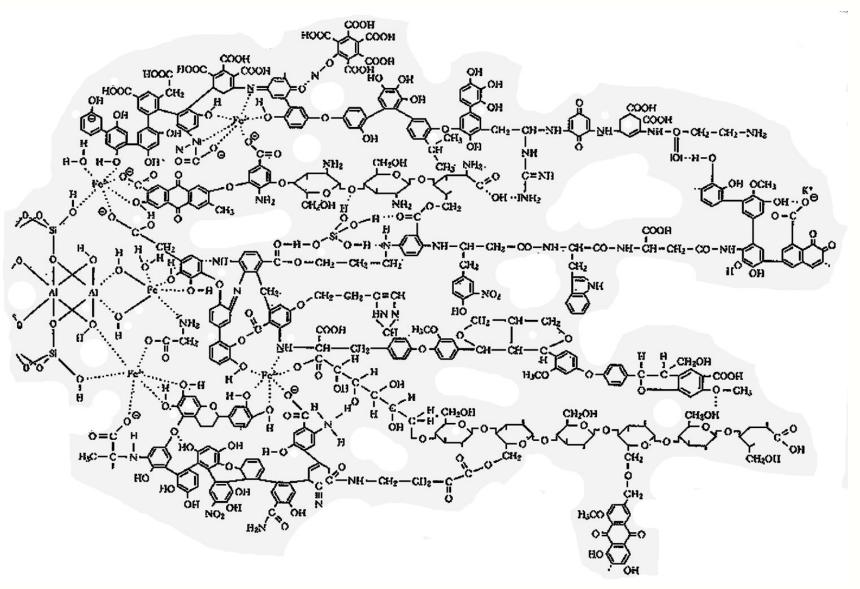
Structure Attributed to Gamble et al. (1985)



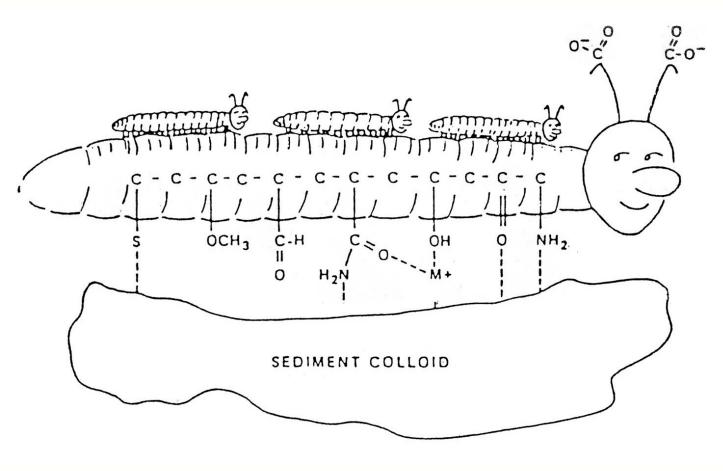
Morel & Hering (1993) Based on Aiken et al. (1985)



Possible Structural Units Set Forth by Averett, Leenheer, McKnight & Thorn (1989) From Morel & Hering, 1993



Kleinhempel reprinted from Albrecht Thaer Archiv (1970)



Organic Solute Macromolecule (ORSMAC) Leenheer 1985) Molecular model of the lowest energy conformation of humic acid building blocks

Carbon atoms-green Oxygen atoms-red Nitrogen-blue Hydrogen not shown

Davies & Ghabbour, 1999

Importance of Humic Materials

Global Carbon Reservoir Take Part in Interfacial Phenomena Undergo Coagulation and Aggregation Involved in Photochemical Reactions **Contain Radicals Known Reducing Agents Methylate Metals** Form Chlorinated Species, THMs DBPs **Detoxify Metals Limit Bioavailability of Metals Alter Solubility** Influence **Bind Metals & Organic Pollutants Terminal Electron Transport Acceptor for Bacteria**

The Removal of Dissolved Humic Acid During Estuarine Mixing

L. E. Fox^a

College of Marine Studies, University of Delaware, Lewes, Delaware, USA Received 2 February 1982 and in revised form 28 May 1982

Keywords: humic acids; dissolved organic compounds; estuaries

A simple method for the determination of dissolved humic acid based on carbon analysis is presented. This method was used to measure the distribution of dissolved humic acids in seven coastal plain estuaries located in the middle-Atlantic United States. Results indicate that 100% of the dissolved humic acid was removed during estuarine mixing, although concurrent measurements of dissolved organic carbon showed either production or conservative behavior in regions of the estuary where humic acid removal was observed. It is apparent from these observations that removal of dissolved humic acid is a minor part of the estuarine transport of dissolved organic carbon.

Laboratory experiments carried out by mixing river water with sea water demonstrated that salt-induced removal of dissolved humic acid was insignificant in two of three estuaries studied. These results suggest *in situ* removal of dissolved humic acid may not be universally caused by increasing estuarine salinity.

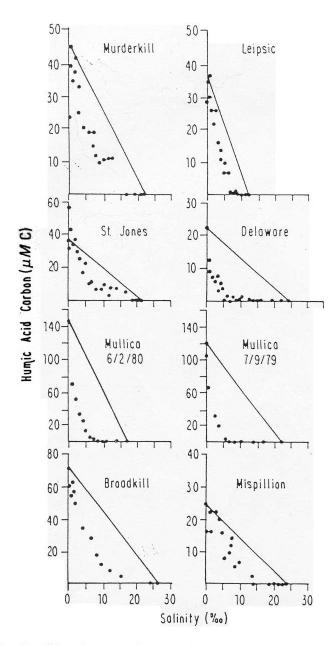




Figure 3. Humic acid carbon as a function of salinity. The standard error of the mean is 5%.

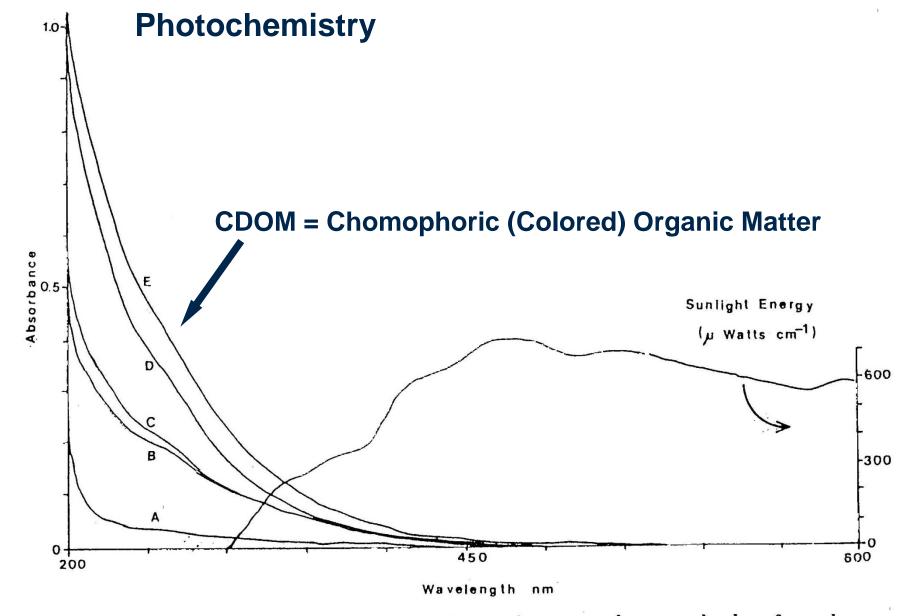
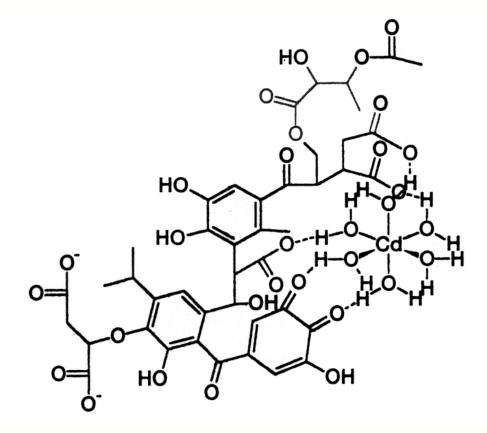
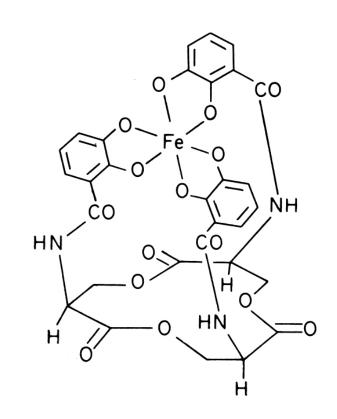


Figure 1. Absorption spectra (pathlength of 1 cm) of several waters and a generalized surface solar energy distribution (adapted from ref. 8). (DOC of waters: A = 3.0, B = 7.8, C = 13.4, D = 13.4, $E = 15.4 \text{ mg } L^{-1}$).

Metal Complexation by Humic Materials





Leenheer et al. (1998)

Morel (1983) 36

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- Fox, L. (1983) Estuarine Coastal Shelf Sci. 16, 431
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Chemical Oceanography Organics III

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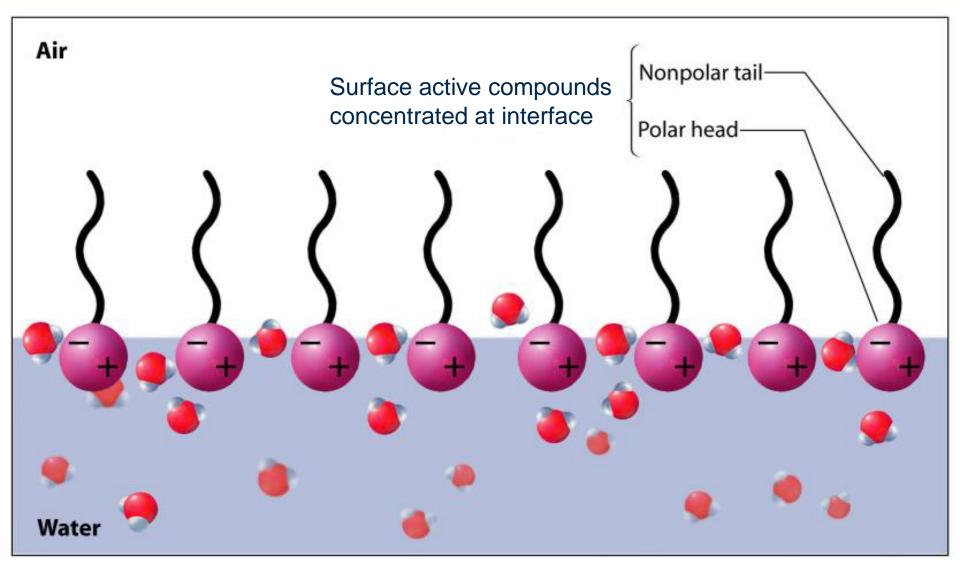
http://faculty.uml.edu/David_Ryan/84.653



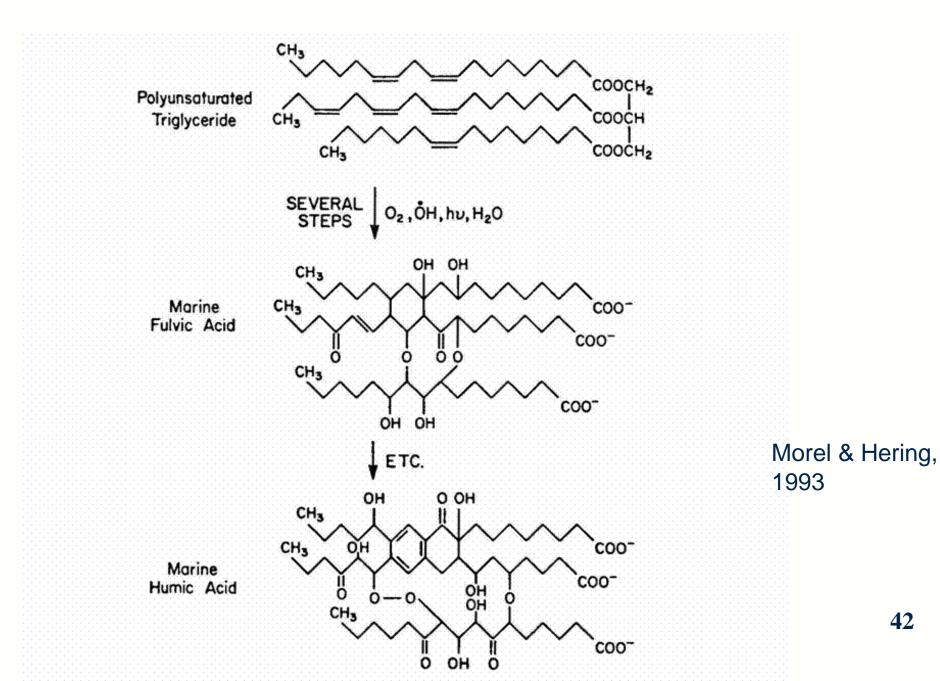
Sea Foam caused by naturally occuring surface active agents



Air-Sea Interfacial Chemistry



From Conceptual Chemistry, Second Edition by John Suchocki. Copyright © 2004 Benjamin Cummings, a division of Pearson Education.



Importance of Humic Materials

Global Carbon Reservoir Take Part in Interfacial Phenomena Undergo Coagulation and Aggregation **Involved in Photochemical Reactions Contain Radicals Known Reducing Agents Methylate Metals** Form Chlorinated Species, THMs DBPs **Detoxify Metals Limit Bioavailability of Metals** Alter Solubility Influence **Bind Metals & Organic Pollutants Terminal Electron Transport Acceptor for Bacteria**

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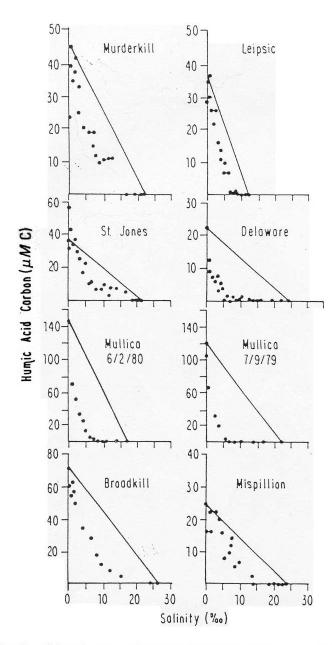
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Fox, 1983

Figure 3. Humic acid carbon as a function of salinity. The standard error of the mean is 5%.

TABLE 10.2Photoreactions of Organic CompoundsChromophoreProducts or effects

| Humic, fulvic | 1. Bleaching of absorption and fluorescence | | | | |
|---|---|--|--|--|--|
| | 2. Production of singlet oxygen | | | | |
| | 3. Fe(III) reduction | | | | |
| | 4. Release of soluble P | | | | |
| | 5. Oxidation of cumene via ROO and OH radicals | | | | |
| | 6. Oxidation of phenolic groups to ArO and formation of e^- and O_2^- | | | | |
| | 7. CO formation | | | | |
| | 8. H_2O_2 formation (via O_2^- ?) | | | | |
| Chlorophyll | Loss of chlorophyll | | | | |
| Vitamins | Loss of bioassay activity | | | | |
| Amino acids | ? | | | | |
| Glycine | COOH C-14 loss, HCHO 1 formation | | | | |
| CH ₃ SSCH ₃ CH ₃ S | CH ₃ S | | | | |
| CH ₃ ICH ₃ | CH ₃ Milloro 1006 | | | | |
| Fatty acids | Particles, absorb., hydroperoxides Millero, 1996 | | | | |
| Aldehydes | RCO, R, CO | | | | |

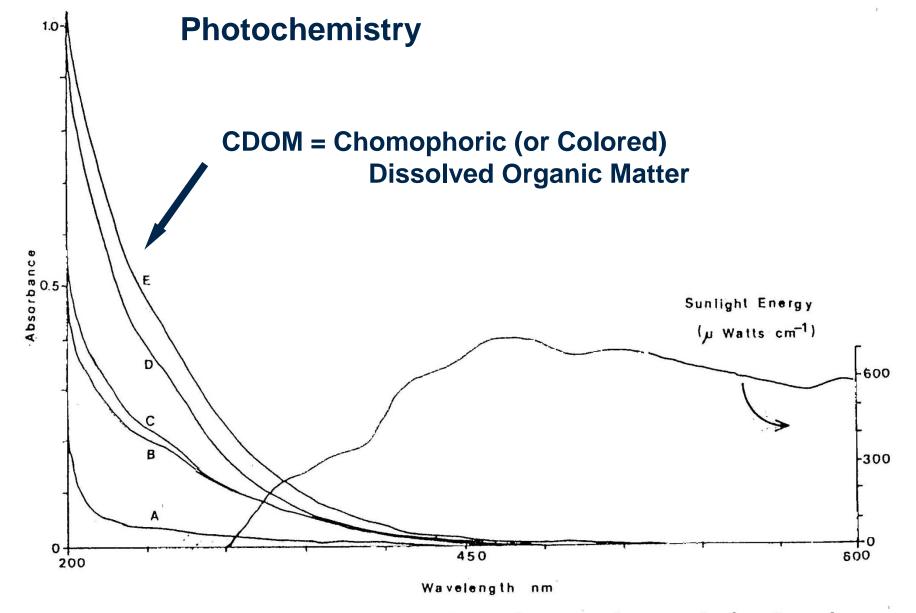
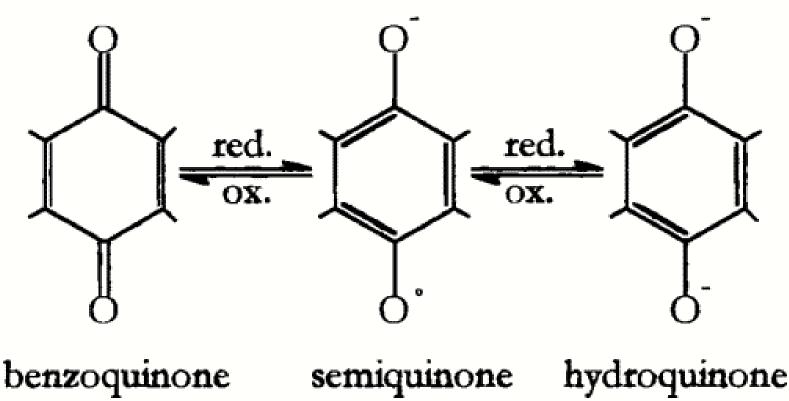


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Quinone radical present in humic material



Scott, McKnight, Blunt-Harris, Kolesar & Lovely (1998) Environ. Sci. Technol. 32, 19

Table 8.3.Representative bulk methods of organic characterizationMALDI, matrix-assisted laser desorption ionization; MS, mass spectrometry; CE, capillaryelectrophoresis.

| Analytical method | Measured characteristics | Total parameters | Typical preparation | Required sample |
|--|--|-----------------------------|--|--|
| Elemental Stable isotope Radioisotope Infrared spectra | C, H, O, N, S 13 C, 2 H, 16 O, 15 N, 36 S Δ^3 H and 14 C, Functional groups | 5 5 2 ~20 | ^a Combustion ^a Combustion ^a Combustion ^b Demineralization | <l mg<br=""><l mg<br=""><l mg<br=""><l mg<="" td=""></l></l></l></l> |
| NMR spectra Pyrolysis-MS MALDI-MS CE-MS | C, H, N and P types Degradation products Intact molecules Intact molecules | ~30 >100 >100 >100 | ^b Demineralization ^b Demineralization ^b Demineralization ^b Demineralization | I−10 mg <10 g <1 mg <1 mg |

^{*a*} Combustion is followed by reduction in the analysis of N and H, whereas O₂ is usually generated by pyrolysis.

^bDemineralization includes separation from mineral phase by organic extraction or mineral dissolution and is typically necessary for sediments but not pure organic materials.