	den site dan site den substantion de la sector de				
PROSJEKTRAPPORT			Distribusjon: ÅPEN		
			AFEN		
	A	ISSN 0071-5638	HI-prosjektnr.:		
			92.01.03		
			Oppdragsgiver(e):		
HAVFORSK					
	en 2 Postboks 1870 50 23 85 00 Faks: 55 23				
Forskningsstasjonen Flødevigen 4817 His Tlf.: 37 05 90 00 Faks: 37 05 90 01	Austevoll Havbruksstasjon 5392 Storebø Tlf.: 56 18 03 42 Faks: 56 18 03 98	Matre Havbruksstasjon 5198 Matredal Tlf.: 56 36 60 40 Faks: 56 36 61 43	Oppdragsgivers referanse:		
Rapport:					
FISKEN OG HA	<b>AVET</b>		NR. 18 - 1998		
Tittel:			Senter:		
A USER'S GUIDE TO	A USER'S GUIDE TO NORWECOM V2.0.				
THE NORWEGIAN ECOLOGICAL MODEL SYSTEM			Seksjon:		
			Fysisk oseanografi		
Forfatter(e):		<u></u>	Antall sider, vedlegg inkl.:		
Morten D. Skogen og Henrik Søiland			42		
			Dato:		
			18.12.98		
Sammendrag:					
A coupled 3 dimensional Physical-Chemical-Biological model system, NORWECOM, has been developed through cooperation between several Norwegian institutions. The physical module is based on the Princeton Ocean Model already well documented in several reports. This guide is an attempt to give an introduction to the Chemical-Biological module, and will hopefully help new users to understand the flow between and processes inside the various subroutines. The guide will concentrate on how the chemical-biological					

dynamics is formulated in the module. Why these formulations is preferred to others, is not a part of such a guide.

Emneord - norsk: 1. Numerisk modell 2. Bruker manual

Hundbyik Prosjektleder

Emneord - engelsk: 1. Numerical model 2. User's guide

Einas Semusican Seksjonsleder

-

### A User's guide to

# NORWECOM v2.0

The NORWegian ECOlogical Model system

Morten Dahlberg Skogen and Henrik Søiland

Institue of Marine Research Division of Marine Environment P.O.Box 1870 N-5024 Bergen-Nordnes morten@imr.no henrik@imr.no

December 1998

\*\*\*\*\*

\*

. .

#### Foreword

A coupled 3 dimensional Physical-Chemical-Biological model system, NORWECOM, has been developed through cooperation between several Norwegian institutions. The physical module is based on the Princeton Ocean Model already well documented in several reports. This guide is an attempt to give an introduction to the Chemical-Biological module, and will hopefully help new users to understand the flow between and processes inside the various subroutines. The guide will concentrate on how the chemical-biological dynamics is formulated in the module. Why these formulation is preferred to others, is not a part of such a guide.

-

# Contents

1	Intro 1.1	oducti What's	on s new ?	<b>1</b> 2			
2	<b>The</b> 2.1	e Physical Model The main processes					
3	$\mathbf{The}$	The Chemical-Biological Model					
	3.1		ain processes	7			
		3.1.1	Incident irradiation	8			
		3.1.2	Light in the water column	11			
		3.1.3	Primary production	12			
		3.1.4	Nutrients	13			
		3.1.5	Oxygen	13			
		3.1.6	Sedimentation and resuspension	13			
	3.2	The di	fferential equations	15			
	3.3 The subroutines		16				
		3.3.1	Light routines	16			
		3.3.2	Biochemical routines	17			
		3.3.3	Transport routines	18			
		3.3.4	Include files	20			
	3.4	The C	hemical-Biological variables	20			
	3.5		upplied input/initialization	24			
4	Same		al negulta and examples	<b>27</b>			
4	Som	ie mod	el results and examples	41			
A	The	includ	le file BIOCOM	33			
в	The	includ	le file SEDCOM	35			

i

CONTENTS

ii

# Chapter 1 Introduction

NORWECOM, the NORWegian ECOlogical Model system, is a coupled three dimensional physical, chemical and biological model system applied to study primary production and dispersion of particles (fish larvae, pollution). The model is the result of a cooperation between several Norwegian institutions. The coupled physical-chemical-biological modeling activity of the whole North Sea started in 1990-91 through close cooperation between the Institute of Marine Research (IMR) in Bergen, the Department of Fisheries and Marine Biology (IFM) at the University of Bergen, and The Norwegian Meteorological Institute (DNMI) in Oslo. Later, also Department of Mathematics at the University of Bergen has taken part in the development of NORWECOM.

The physical module is based on the well known Princeton Ocean Model (POM) by Blumberg & Mellor (1987). This model has been applied successfully in many regions after its first version was presented in 1977 (Blumberg & Mellor, 1980). The model also performed favorably in a Norwegian model evaluation project, MOMOP (Martinsen *et al.*, 1990; Slørdal *et al.*, 1991). It is a clear strength that the chemical-biological module is run coupled to a well known physical model, instead of using more or less realistic dispersion techniques.

The chemical-biological module is based on a paper by Aksnes & Lie (1990) who used it in a land-locked fjord, Lindåspollene, in western Norway. After this the model has been further developed and studied, and implemented on a model area covering an extended North Sea. This implementation was used by Aksnes *et al.* (1995) for simulating the *Chrysochromulina polylepis* bloom in Skagerrak in 1988. In the same paper the nutrients and phytoplankton development in six enclosures were simulated in order to do a verification of the validity of the set of parameters proposed by Andersen & Nival (1989). Skogen *et al.* (1995) used NORWECOM for simulating the time variability of both nutrient distributions and primary production in 1985 in the whole North Sea. In the same paper they also studied the possible effects of reducing the anthropogenic riverine loads and atmospheric nutrient deposition, and the transport of nutrients to the Skagerrak.

The main weakness of 3-D modeling activities claiming to simulate nature is the lack of comparison with adequate real data. Using the extensive SKAGEX dataset (Danielssen *et al.*, 1991, 1997; Ostrowski, 1994) the model has been validated (Dee, 1994) in Svendsen et al. (1996a); Skogen et al. (1997c). Additional model validation through comparison with field data can also be found in Aksnes et al. (1994); Berntsen et al. (1996); Svendsen et al. (1995c); Søiland & Skogen (1998). The model has been used to investigate long-term residual transports in the North Sea together with two other models (Smith et al., 1996), and taken part in several model to model intercomparison studies (Røed et al., 1989; Gustafsson & Jönsson, 1995; Proctor et al., 1997; OSPAR, 1998).

In recent years, the model has been used in new ares and for a wide range of applications. In Laane *et al.* (1996); Skogen *et al.* (1997b, 1998); Skogen & Moll (1998); Skogen (1998b); Svendsen *et al.* (1995a,b); Søiland *et al.* (1996) the model has been used for estimation of transports and primary production, and investigation of eutrophication in the North Sea and Skagerrak. In Iversen *et al.* (1998); Skogen *et al.* (1997a); Svendsen *et al.* (1995c, 1996b); Sætre *et al.* (1998); Walsh *et al.* (1996) the model, coupled with a Lagrangian transport model, has been used in studies of fish recruitment and assessment, and in Skogen (1998a) a version of NORWECOM has been implemented for the Benguela upwelling system.

The model is described in parts in several of the above mentioned references. The first thorough presentation of the model, including data flow and the subroutines, was given in *A Users guide to NORWECOM* (Skogen, 1993). The present report is an updated version of that user's guide, including the extensions and changes that has been done based on the experiences the last years. This guide will only discuss how things are done within the model, to answer questions like why one formulation is preferred to another, is not a part. This report can be downloaded in postscript format from the NORWECOM home page http://www.ii.uib.no/~morten/norwecom.html

### 1.1 What's new ?

The main change from NORWECOM version 1.0 to NORWECOM version 2.0 is the inclusion of oxygen as a prognostic variable, and the inclusion of a sedimentation and resuspension module. A major limitation of the previous version was also the lack of regeneration of silicate, due to this the model could only be run one year at a time. In this version a new prognostic variable, biogenic silica (diatom frustules), is included, and the model can therefore be run for more than one year, without reinitialization.

In addition there has been a change in a few parameters, and several updates in the physical formulation and forcing.

The content of this guide is very similar to the previous one (Skogen, 1993). However, two extensions are done. A tabular showing the value of all constants given in Section 3.4, and one extra chapter is included to show some examples and validation results from several different model runs.

# Chapter 2

# The Physical Model

The circulation model is based on approximations from the three-dimensional, primitive equation, time-dependent, wind and density driven Princeton Ocean Model (POM). The model is fully described in Blumberg & Mellor (1987); Mellor (1996); O'Connor (1991). Therefore only a short review of the main physical processes is presented here. For a more thorough study of the physical processes, see the references above, or find more information at the POM homepage at : http://www.aos.Princeton.EDU/WWWPUBLIC/htdocs.pom/

### 2.1 The main processes

The prognostic variables of the Princeton Ocean Model are :

- three components of the velocity fields
- temperature
- salinity
- turbulent kinetic energy
- turbulent macroscale
- water level

The governing equations of the model are the horizontal momentum equations, the hydrostatic approximation, the continuity equation, conservation equations for temperature and salinity and a turbulence closure model for calculating the two turbulence variables (Mellor & Yamada, 1982). The equations and boundary conditions are approximated by finite difference techniques. In the vertical a sigma-coordinate representation is used, and the horizontal grid uses cartesian coordinates in a staggered *Arakawa C grid* (Mesinger & Arakawa, 1976).

The physical forcing factors are atmospheric wind and pressure, tides, freshwater runoff, surface heat fluxes and in- and out-flows at the open boundaries. The freshwater runoff is handled by routines developed by Berntsen and Ådlandsvik. Here, the water level is increased in the grid point nearest to the river outlet, and the salinity updated assuming a homogeneous upper layer of 3 meter. Special routines for the exchange between external

basins (e.g. the Baltic) and the model area are included and based on techniques due to Stigebrandt (1980). In the lack of data on the surface heat fluxes, a "relaxation towards climatology" method is used (Cox & Bryan, 1984). During calm wind conditions, the surface temperature field will adjust to the climatological values after about 10 days (Oey & Chen, 1992). The net evaporation precipitation flux is set to zero.

Around the open boundary a Flow Relaxation Scheme (FRS-zone) is used (Martinsen & Engedahl, 1987). Each prognostic variable,  $\phi$ , in the zone is simply updated by the translation  $\phi = (1 - \beta) \phi_{int} + \beta \phi_{ext}$ , where  $\phi_{int}$  contains the time integrated, unrelaxed values calculated in the entire model domain, i.e. also in the areas covered by the FRS-zone, and  $\phi_{ext}$  is the specified external solution in the zone.  $\phi$  is the new value and  $\beta$  a relaxation parameter which varies from 0, at the end of the zone facing the interior model domain, to 1, at the outer end of the zone. This simple FRS zone technique effectively absorbs inconsistencies between forced boundary conditions and model results.

For modeling in the North Sea and Norwegian Sea areas, the model is using six-hourly hindcast atmospheric pressure fields provided by the Norwegian Meteorological Institute (DNMI) (Eide *et al.*, 1985), 6-hourly wind stress (translated from the pressure fields by assuming neutral air-sea stability) and four tidal constituents (M2,S2,K1,O1). Initial values for velocities, water elevation, temperature and salinity are taken from monthly climatologies (Martinsen *et al.*, 1992). Interpolation between these monthly fields are also used at all open boundaries.

In the POM-model the horizontal time differencing is explicit whereas the vertical differencing is implicit. The latter eliminates time constraints for the vertical coordinates and permits the use of fine vertical resolution in the surface and bottom boundary layers. The model has a free surface and a split time step. The external mode portion of the model is two-dimensional and uses a short time step based on the CFL (Courant Friedrichs Levy) requirement related to the *external* wave speed (the fastest wave), while the internal mode is three dimensional and uses a much longer time step based on the same CFL requirement, but now related to the *internal* wave speed.

There are many different constants to be initialized in the physical module. Two of them need special attention, since the physical results are highly dependent on their values (Berntsen *et al.*, 1996).

UMOL  $(m^2/s)$ , the minimum allowed value for the vertical eddy viscosity/diffusion coefficient (which are calculated in connection with the turbulence closure model), causes vertical gradients to be smoothed, and should therefore be kept at a minimum. Unfortunately, too small values may cause instabilities. To avoid this, three different UMOLs are introduced in NORWECOM. One is related to the vertical gradients in horizontal velocity, one for the vertical temperature and salinity gradients and one for the chemical/biological gradients. The first one is given the default value set in POM of  $2 \cdot 10^{-5}$ , while the two others are set to  $10^{-7}$ .

By comparing with data there are clear indications that using the default value of  $2 \cdot 10^{-5}$  is smoothing the vertical salinity gradients too much in strongly stratified waters. Using  $10^{-7}$  for the minimum temperature and salinity vertical diffusion coefficient, seems to give much more realistic results. However, to avoid instability problems which occurred

#### 2.1. THE MAIN PROCESSES

in the most shallow and stratified waters with this value, a simple filtering and surface boundary condition procedure has been added.

HORCON is a dimensionless horizontal diffusion constant used in connection with the Smagorinsky diffusivity for horizontal diffusion (Smagorinsky, 1963),

$$A_M = HORCON \cdot \Delta x \ \Delta y \ \frac{1}{2} \left[ \left( \frac{\partial u}{\partial x} \right)^2 + \frac{1}{2} \ \left( \frac{\partial v}{\partial x} + \frac{\partial u}{\partial y} \right)^2 + \left( \frac{\partial v}{\partial y} \right)^2 \right]^{\frac{1}{2}}.$$

The technique is used for eliminating small scale  $(2\Delta x)$  velocity-and density structures being produced in the model (and in nature) but not being resolved. A large value for HOR-CON increases the horizontal diffusion. If the grid spacing is small enough, HORCON can be null. In the model a value of 0.20 is used for the velocities and the chemical/biological variables, while 0.03 is used for temperature and salinity.

CHAPTER 2. THE PHYSICAL MODEL

# Chapter 3

# The Chemical-Biological Model

The first version of the biological-chemical module was first described Aksnes & Lie (1990). For further applications and model description see also Aksnes *et al.* (1995); Skogen *et al.* (1995).

This chapter describes the main processes, the differential equations, the different subroutines and a list of the chemical and the biological variables and constants as used in NORWECOM v2.0.

### 3.1 The main processes

The biological model is coupled to the physical model through the subsurface light, the hydrography and the horizontal and the vertical flow and mixing of the water masses. NORWECOM differs between two groups of phytoplankton, diatoms (DIA) and flagellates (FLA), and their growth is affected by the nutrient concentration, light intensity and temperature. The nutrients are represented by inorganic nitrogen (NIT) such as nitrate and ammonia, inorganic phosphorus (PHO) (phosphate) and inorganic silicon (SIL) (silicate). The main difference between diatoms and flagellates, is that silicate is not rate limiting for the production of flagellates. The nutrients are added to the system through the river runoff and from the atmosphere, further nitrogen and phosphorus are regenerated from the dead algae, detritus (DET), and silicate from biogenic silica (SIS) at a constant rate. Oxygen (OXY) is released during primary production (growth) and it is consumed in respiration and when detrital matter is regenerated. Oxygen is assumed to be saturated at the surface.

For both the dead and living algae a sinking rate which may depend on the nutrient concentrations, is included. There are no zooplankton eating the algae. Instead a constant death rate, that also should include grazing, is included. Thus in this context, also zooplankton is included as a forcing variable. The particles (DIA, FLA, DET and SIS) may settle at the bottom, and can either be buried in the sediments or resuspended. The sedimentation and resuspension is a function of the bottom stress, which in the model is the sum of the stress due to surface waves, and the bottom currents, including tides. Algae that settle at the bottom is transformed immediately to detritus. Detritus mineralize in the sediment at the same rate as in the pelagic, but a constant fraction of the remineralized nitrogen is lost due to denitrification. Processes like aggregation and flocullation affect the sedimentation and resuspension rates, but are not included. A limitation is also the lack of realistic light attenuation due to riverine inputs of yellow substance etc..

There is a total of ten prognostic variables in the chemical-biological part. In addition to the eight already mentioned, NIT, PHO, SIL, DET, SIS, FLA, DIA and OXY, also light in the water column, RAD, and an inorganic suspended particulate matter variable ISPM, are included. The sinking speed used for ISPM is tuned towards small particles (< 50  $\mu$ m). All variables are defined at the modeled temperature and salinity, (S,T), points of the Arakawa-C grid. All chemical-biological concentrations in the model are given in  $(mg/m^3)$ s, except ISPM which is given as (g/l). A schematical overview of the different processes is given in Figure 3.1.

#### 3.1.1 Incident irradiation

The incident irradiation is modeled using a climatological light formulation (Skartveit & Olseth, 1986, 1987), that have been updated to include real light measurements (Ulvestad, pers.comm). In this section, first the climatological formulation will be presented, and than the updated real-light algorithm will be given.

The original formulation gives a *mean* irradiation as a function of latitude and season. This mean climatological irradiance, varies according to a presumed sinusoidal variation, and all constants,  $a_x - f_x$ , should be specific for the area to be modeled. The irradiance is split into a diffuse (x = dif) and a direct (x = dir), (or beam) component

$$H_x(h,n) = I_0(n) \cdot Tr_{0x}(n) \cdot F_x(h).$$
(3.1)

Here  $H_x(h, n)$  is either direct or diffuse irradiance at the surface,  $I_0(n)$  the solar irradiance at normal incidence just outside the atmosphere,  $Tr_{0x}(n)$  the transmittance and  $F_x(h)$ the solar elevation function. Further h is the solar elevation and n the day number. The transmittance at overhead zenith sun is given by :

$$Tr_{0x}(n) = a_x(1 + b_x \cos\frac{n - c_x}{365}2\pi).$$
(3.2)

The solar elevation function,  $F_x(h)$ , expresses the effect of varying solar elevation, which is estimated in every internal time step, and is given by :

$$F_x(h) = d_x + e_x \sin h - f_x (\sin h)^{1/2}.$$
(3.3)

The formulas are valid when the solar elevation is above 5 degrees, nevertheless they are used for all solar elevations.

In case of large day to day variability in the incoming light, such a climatological approach is not good enough if one wants to initiate or simulate special blooms. Therefore an interpolation technique for the parameters  $a_x - f_x$ , in the case when real light measurements are available, has been proposed.

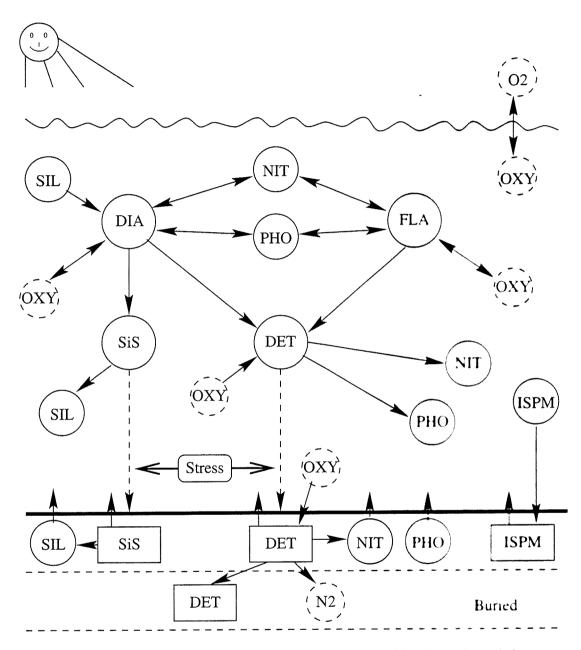


Figure 3.1: Schematical overview of the chemical-biological module

As input the global irradiance per day is needed for one or several distinct points in the model domain. One must also define the (sub)domains where these time series are assumed valid. The constants  $a_x \ldots f_x$  in (3.2) and (3.3) depends on the light intensity, thus in their papers Skartveit & Olseth differ between clear sky and all days average constants. Given these values for global, diffuse and direct light, the new algorithm interpolates and incorporates the parameters  $a_x \ldots f_x$  in the following way.

Given one point, a domain and a time series of global daily irradiance :

```
main loop
С
C For each day find the transmittance (3.2)
C
   if (newday)
      meanglo = global irradiance (daynr, average constants)
      maxglo = global irradiance (daynr,clear sky constants)
      read (irrad_file), global = global irradiance (daynr)
      if (global < meanglo)
          lowirrad = .true.
      else if (global > maxglo)
          compute transmittance from clear-sky-constants
      else
          interpolate new constants from average and clear sky
          compute transmittance using interpolated constants
      end if
   end if
С
C At every time step find direct and diffuse irradiance (3.1)
С
   find sun height (longit, latit, daynr, time)
   if (lowirrad)
      direct = 0.
      diffuse = global(sunheight,daynr,aver_const.)
   else if (global > maxglo)
      compute the irradiances from clear-sky-constants and sun height
   else
      interpolate new constants from average and clear sky
      compute the irradiances using interpolated constants and sun height
   end if
С
С
   continue (main loop)
```

10

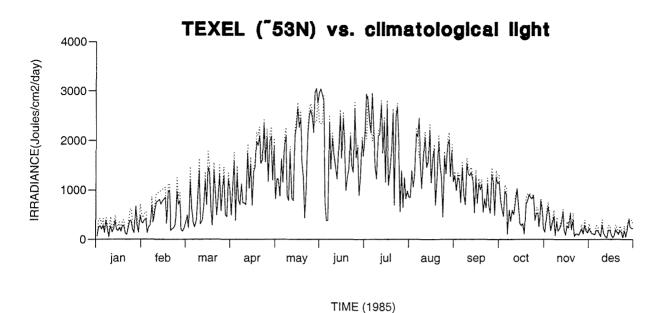


Figure 3.2: Modeled global irradiance (dotted line) vs. real measurements at Texel (53  $^o\!N)$  in 1985

As seen it is now differed between three cases. In the first one with high global irradiance, the *clear-sky* constants are used, and in the second one with medium light, a set of *interpolated* constants are defined. In the third one with low irradiance, the *average* constants are used, but the light formulations are changed (see Equation (8) in Skartveit & Olseth (1986)) such that the direct irradiance is set equal to zero and the diffuse irradiance is set equal to a global hourly irradiance function, which for hour j and day n is given as :

$$\bar{H}_{x}(h_{j},n) = \bar{H}_{x}(n) \cdot \frac{F_{x}(h_{j})}{\sum_{i} F_{x}(h_{i})},$$
(3.4)

where  $\bar{H}_x(n)$  is the given global irradiance for the actual day, and  $F_x(h)$  is the solar elevation function.

This new real-light formulation enables the mapping of the day to day variations in the global irradiance in a very precise way. In Figure 3.2 a comparison between the modeled and measured light at Texel  $(53 \, {}^{o}N)$  in 1985 is given.

#### 3.1.2 Light in the water column

The diffuse light is calculated from

$$I_{dif}(x, y, z, t) = PAR \cdot R_{dif}(x, y, t) e^{-\frac{\kappa(x, y, z, t)}{\mu}},$$
(3.5)

where  $R_{dif}(x, y, t) = H_{dif}(h, n)$ , the diffuse component of the surface irradiance, and *PAR*, photosynthetic available radiance, a constant which converts from incident diffuse irradiation to photosynthetic active irradiation.  $\mu$  is the mean cosine of the diffuse light (Platt & Jassby, 1976), and  $\kappa$  the attenuation coefficient which is kept as a function of the concentration of chlorophyll and "other substances".

$$\kappa = b_2 z + \frac{\nu}{N2Chla} \int_0^z (DIA(x, y, z, t) + FLA(x, y, z, t)) dz.$$
(3.6)

Here  $\nu$  is the chlorophyll-a light extinction coefficient, N2Chla the fraction of nitrate and chlorophyll\_a in a cell, and  $b_2$  extinction due to water and other substances.

A similar formulation to (3.5) is given for the direct light,  $I_{dir}(x, y, z, t)$ , by substituting  $R_{dif}$  with  $R_{dir}$  and  $\mu$  with  $\cos \phi$ , where  $\phi$  is the zenith angle of the direct light in the water column computed from Snells formula,  $n_1 \cos h = n_2 \sin \phi$ , where  $n_1$  and  $n_2$  are the refraction coefficients for air and water.

#### 3.1.3 Primary production

The concentration of chlorophyll in the system is affected by the production of the algae, their death rate and respiration. Since zooplankton is not included in the model, the death rate of algae also includes possible grazing.

The relationship between phytoplankton production and light intensity, and the relationship between phytoplankton production and nutrient uptake is represented by an affinity formulation, see Aksnes *et al.* (1995). The combined effects of nutrient and light limitation are multiplicative and given by :

$$\mu_{dia}(x, y, z, t) = \mu_{max} \cdot V_1 \cdot N_{lim} \cdot Dia(x, y, z, t), \quad N_{lim} = \min_{2 \le i \le 4} V_i, \quad (3.7)$$

where

$$V_{i} = \frac{S_{i}}{S_{i} + \frac{\mu_{max}(T)}{\alpha_{i}}}, \ i = 1, \dots, 4$$
(3.8)

is a modified Michaelis-Menten limitation for substance  $S_i$ . In the equations i = 1 corresponds to irradiance, i = 2 to nitrate, i = 3 to phosphate and i = 4 to silicate. In this formulation the constant half saturation parameters,  $K_s$ , are avoided. Instead they are made temperature dependent through the affinity parameter,  $\alpha_i$ , defined as :

$$\alpha_i = \frac{\mu_{max}(T_0)}{K_-Dia_i},\tag{3.9}$$

where  $K_Dia_i$  is the conventional half saturation constant for diatoms for substance number i at temperature  $T_0$ .  $\mu_{max}$  is the specific growth rate of the population under optimum light and nutrient conditions and is made temperature dependent as suggested in Eppley (1972). The relation

$$\mu_{max}(x, y, z, t) = a_1 e^{a_2 T(x, y, z, t)}, \qquad (3.10)$$

#### 3.1. THE MAIN PROCESSES

is chosen, where  $a_1$  is the diatom production maximum at  $0 \ C$   $(s^{-1})$ , and  $a_2$  is the production rate temperature dependence  $(C^{-1})$  for diatoms.

The respiration is assumed to be related to the temperature according to the equation

$$R_{dia}(x, y, z, t) = a_5 Dia(x, y, z, t) e^{a_6 T(x, y, z, t)},$$
(3.11)

with  $a_5$  and  $a_6$  the repiration rate at  $0 \mathcal{C}(s^{-1})$  and the respiration rate temperature dependence  $(\mathcal{C}^{-1})$  respectively.

The death rate of the algae (in the whole water column) is assumed to be constant as long as the chlorophyll concentration of the algae at the surface is above a minimum level, and zero if it is below. This is because the biological model is light limited in winter, and in order to prevent the algae in the model to extinct, further death when their concentration becomes too small, must be prevented.

All these expressions refers to the diatoms. Analogous formulations are used for the production of flagellates. The only difference is that silicate is not rate-limiting for the flagellates.

#### 3.1.4 Nutrients

The primary production is limited by three different nutrients, NIT, PHO and SIL<sup>1</sup>, which are transported together with the water masses, and are added to the system through river input, atmospheric deposition and by regeneration of NIT and PHO from dead organic matter (DET) and SIL from the biogenic silica (SIS). Inside the algae, both dead and alive, PHO and SIL are expressed as NIT-equivalences, i.e. the uptake of PHO and SIL through the photosynthesis is expressed as a constant times the uptake of NIT. The Redfield ratios are used for these cellular ratios. For more details on the uptake and regeneration see the differential equations in Section 3.2.

#### 3.1.5 Oxygen

The oxygen concentration is affected by the primary production, respiration and remineralization of detrital matter. The amount of oxygen released by primary production is proportional to the amount of inorganic nitrogen consumed, and is given by constant (Redfield ratio). The same ratio for oxygen consumption is used for the respiration and remineralization process. The ratio is based on the assumption that inorganic nitrogen is converted from nitrate to organic matter and vice versa. For the fraction of nitrogen that is denitrified in the sediments a somewhat smaller consumption takes place.

#### 3.1.6 Sedimentation and resuspension

Particulate matter (ISPM) has a sinking speed relative to the ambient water and thus may reach the bottom. However laboratory measurements, and also field experience, show

<sup>&</sup>lt;sup>1</sup>Flagellate production is not limited by SIL

that if the bottom stress is above a threshold value it does not settle on the bottom, but remains suspended. If the stress is below a certain value, particles settle on the bottom. The opposite process of resuspension is also dependent on the bottom stress, such that resuspension occurs if the stress is above a certain limit. The value at which resuspension occurs is higher than the maximum stress value for sedimentation to occur, such that there is a small window at which neither sedimentation nor resuspension takes place. These limits are in general a function of the properties of the particles (sinking speed), and also of the time since sedimentation occurred (armouring). Since the values are variable, quite a scatter of values for these limits are occur in the literature, but we have chosen values used in other North Sea models (Pohlmann & Puls, 1994; Mayer, 1995). The sedimentation and resuspension are controlled by the following relationships :

$$\begin{aligned} \tau &< \tau_1, \qquad \Rightarrow sedimentation \\ \tau_1 &< \tau &< \tau_2, \\ \tau_2 &< \tau \qquad \Rightarrow resuspension, \end{aligned}$$
 (3.12)

where  $\tau_1 = 0.064 \ Pa$  and  $\tau_2 = 0.78 \ Pa$ . In a shallow area such as the North Sea the bottom stress is due to both the current and surface waves. In NORWECOM the total bottom stress,  $\tau_B$ , is calculated as the sum of these two components.

$$\tau_B = \tau_{wave} + \tau_{current} . \tag{3.13}$$

The magnitude of the wave stress is given by :

$$\tau_{wave} = \frac{1}{2} \ \rho \cdot f_w \cdot U_w^2,$$

where  $f_w$  is the wave friction factor and  $U_w$  is the wave orbital velocity close to the bottom, while the bottom current stress,  $\tau_c$ , is taken from the physical model. The stress due to surface waves is based on wave data from the WINCH wave model (SWAMP-Group, 1985; Reistad *et al.*, 1988).

The resuspension rate is proportional to the bottom stress and is given by :

$$\epsilon = M (\tau_B - \tau_2), \qquad M = 10^{-4} m^{-1}.$$
 (3.14)

There are several forms of particulate matter in the sediment, and if resuspension occurs they are resuspended in the fraction of the total that they occur.

The algae that sediment out to the bottom is immediately transferred to the detrital pools (DET and SIS (only DIA)). Since algae is transferred to these detrital pools there are only two forms of detrital matter (DET and SIS) and the inorganic particulate matter variable (ISPM) present in the sediment. Here the detrital matter is mineralized back to inorganic nutrients using the same rate constants as in the pelagic. Oxygen is not a variable in the sediments so the oxygen consumed in this process is drawn from the water just above the sediments (lowest sigma layer). A constant fraction of the nitrogen that is remineralized in the sediments may also be transformed to nitrogen gas through denitrification Lohse *et al.* (1996).

Another potential sink term for nutrients in the sediments is the part that is buried. Burying is formulated as a rate term for the amount of detrital matter above a given value. This is a slow process (seasonal) and it does not prevent the high concentrations of detrital matter in the sediments giving rise to unrealistic high fluxes of nutrients from the sediments. To keep the remineralization at a reasonable level a max value for this release based on published values is assumed (Gehlen *et al.*, 1995; Lohse *et al.*, 1995).

### 3.2 The differential equations

The concentrations of the nine prognostic chemical-biological variables, NIT, PHO, SIL, DET, FLA, DIA, SIS, OXY and ISPM, and the different processes they take part in, can be described using the following set of differential equations :

$$\begin{aligned} \frac{\partial N}{\partial t} + adv(N) &= diff(N) + R_{Dia} + R_{Fla} + cc_4 \, Det - (P_{Dia} + P_{Fla}) + \phi(N) \\ \frac{\partial P}{\partial t} + adv(P) &= diff(P) + cc_1(R_{Dia} + R_{Fla} + cc_4 \, Det - (P_{Dia} + P_{Fla})) + \phi(P) \\ \frac{\partial Si}{\partial t} + adv(Si) &= diff(Si) - cc_2 \, P_{Dia} + scc_4 \, Sis + \phi(Si) \\ \frac{\partial Det}{\partial t} + adv(Det) &= diff(Det) + cc_3 \, (Dia + Fla) - cc_4 \, Det + \phi(Det) \\ \frac{\partial Sis}{\partial t} + adv(Sis) &= diff(Sis) + cc_2 \, (R_{Dia} + cc_3 \, Dia) - scc_4 \, Sis + \phi(Sis) \\ \frac{\partial Dia}{\partial t} + adv(Dia) &= diff(Dia) + P_{Dia} - R_{Dia} - cc_3 \, Dia + \phi(Dia) \\ \frac{\partial Fla}{\partial t} + adv(Fla) &= diff(Fla) + P_{Fla} - R_{Fla} - cc_3 \, Fla + \phi(Fla) \\ \frac{\partial Oxy}{\partial t} + adv(Oxy) &= diff(Oxy) + scc_1 \, (P_{Dia} + P_{Fla} - R_{Fla} - cc_4 \, Det) + \phi(Oxy) \\ \frac{\partial Ispm}{\partial t} + adv(Ispm) &= diff(Ispm) + \phi(Ispm) \end{aligned}$$

Here adv are the terms due to advection, diff the terms due to diffusion, P production terms, R respiration terms,  $\phi$  river and atmospheric deposition (fluxes), or sink and source terms due to sedimentation and resuspension,  $cc_1$  intercellular P/N relationship,  $cc_2$  intercellular Si/N relationship,  $cc_3$  phytoplankton death rate,  $cc_4$  regeneration rate of detritus,  $scc_1$  fraction of oxygen and nitrate for each cell produced, and  $scc_4$  the remineralization rate of biogenic silica.

In more details the operators (after a transformation to the  $\sigma$ -coordinate system) can be written as :

$$adv_{\sigma}(F) = \frac{\partial}{\partial x}(FUD) + \frac{\partial}{\partial y}(FVD) + \frac{\partial}{\partial \sigma}(F\omega)$$
 (3.15)

$$diff_{\sigma}(F) = \frac{\partial}{\partial\sigma} \left(\frac{K_H}{D} \frac{\partial F}{\partial\sigma}\right) + \frac{\partial}{\partial x} \left(DA_H \frac{\partial F}{\partial x}\right) + \frac{\partial}{\partial y} \left(DA_H \frac{\partial F}{\partial y}\right), \qquad (3.16)$$

where F is the substance to be integrated, U and V the horizontal velocities in x and y-direction respectively,  $\omega$  the vertical velocity in the sigma coordinate system,  $D \equiv H + \eta$ (bottom depth + water elevation),  $A_H(m^2s^{-1})$  the horizontal diffusivity and  $K_H(m^2s^{-1})$ the vertical diffusivity. In addition the sinking rates of the particulate variables (DIA, FLA, DET, SiS and ISPM) are applied after the above updates.

### 3.3 The subroutines

The main chemical-biological module is coupled to the physical module through the call to one biology routine, BIOLOGY.F90. In this routine all chemical-biological dynamics is handled in 25 different subroutines. These subroutines can be divided into three groups (light, biochemical and transport) in a natural way. Each of these subroutines will be described shortly in the following subsections. For a more detailed description of the various parameters, see also Section 3.4 and the prologue inside each subroutine. A flowchart is given in Figure 3.3, to illustrate the division, the dependencies between the different subroutines, and the order they are called in. To ease the data flow, two include files (BIOCOM.H and SEDCOM.H) are used. The include files are given in Appendices A and B respectively.

Before BIOLOGY is called, the hydrodynamic forcings should be updated. This includes new computations of the velocity field, the water elevation and fields for horizontal diffusion and vertical eddy viscosity. In addition the bottom stress due to the waves and bottom currents, are calculated before BIOLOGY is called.

#### 3.3.1 Light routines

The biological module starts with calculation of the light conditions. There are two entries to this part, either through the routine DAYNUMB (that are called once every day), or through the routine BIOSUN (see Figure 3.3). All together there are 10 light routines. Five of these (daynumb, declin, comprad, trans, lowlight) are only called through the daynumb entry once a day. The main light routine is RADIATION which calculates the radiation (direct and diffuse) in the whole water column as a function of the direct and the diffuse irradiance at the surface and the *water quality*. The other nine subroutines estimate these dependencies (Equation (3.1) and  $b_2$  in Equation (3.6)). Data for the global daily irradiance is also read here, once every day.

- biosun(daynr,theta,longit,latit,sunti,sunhei) Computes the sun height. h, used in (3.3). Sunhei is input to irrsur and snell.
- comprad(iflag,daynr,theta,clirad,dti) Computes the total climatologically integrated global surface irradiance at daynumber using either mean or clear sky constants from Skartveit & Olseth (1986, 1987). The irradiances are used as input to trans.

daynumb(daynr,ndate) Determines the day number given the date.

- declin(daynr,theta) Computes the declination of the sun,  $\theta$ , as a function of the daynr. The declination is the elevation of the sun measured over the north pole given in degrees.  $\theta$  is used as input to the subroutine biosun.
- irrsur(beamtr,difftr,daynr,diffus,direct,global,low,maxglo,meanglo, sumdf, sunhei, ...) Computes the  $I_0(n)$  part of Equation 3.1, and the solar elevation function (3.3), including the interpolation to find the new constants  $d_x$  and  $e_x$ . Finally the multiplication giving the diffuse and direct components of the surface irradiance (3.1) is done. The irradiances, diffus and direct are input to radiation.
- lowlight(daynr,latit,longit,maxglo,meanglo,sumdf,theta,fsm,dti) Determines the irradiance when the total diurnal integrated irradiance (from data) is lower than the value using the mean sky constants (3.4). This irradiance is input to irrsur.
- radiation(direct,diffus,i,j,kappa,phi,dt,dz) The main light routine. Computes both the diffuse (3.5) and the direct components of the radiation, RAD, in a water column determined by (i,j). It also computes the self shading (integral part of (3.6)), and estimates the attenuation coefficient,  $\kappa$ . The radiation is used as input to the biochemical routine, phyto\_growth.
- snell(sunhei) Computes the zenith angel of direct light,  $\phi$ , in the water column using Snells formula.  $\phi$  is input to radiation.
- trans(daynr,beamtr,difftr,global,low,maxglo,meanglo) Determines the beam and diffuse transmittance (3.2), through the interpolation procedure determining the constants  $a_x, \ldots, c_x$ , and the logical variable *low*. The transmittances is input to **irrsur**, and *low* determines if the routine **lowlight** should be called.
- wqual(kappa,s,fsm) Computes the extinction coefficient,  $b_2$  in (3.6), in the water as a function of water type.  $\kappa$  is input to radiation.

### 3.3.2 Biochemical routines

There are eight biochemical routines with **phyto\_growth**, where the respiration (3.11) and primary production (3.7) are calculated, as the main one. The light that was computed in **radiation** is input to **phyto\_growth**. These routines can logically be divided into two, where one group (**atmnit**, **biosource**, **botsource**, **oxysat**) computes different source terms used by the transport routine vertbio.

affin(aa,kndia,knfla,kpdia,kpfla,ksdia,krdia,krfla) Finds the affinity parameters,  $\alpha_i$ , in Equation (3.9), when the half saturation constants and  $T_0$  are given, and initiates the constants,  $a_i$ , used in Equations (3.10) and (3.11). Affin is only called once.

- atmnit(.....) Updating NIT in the surface layer, according to data for atmospheric deposition. Must be called after the transport routine fctf. Used as input to vertbio.
- biosource(icode,source,pdia,pfla,rdia,rfla,ddia,dfla) Give the source terms, i.e. the values for production, death rate and respiration rate in the differential equations, for all chemical and biological variables. The source term is input to vertbio.
- botsource(icode,source,fsm,dz,dt,dti) Computes the source terms for all chemical and biological variables in the sediment layer, and an additional source term in the next to bottom layer, due to leakage (NIT, PHO, SIL) from the sediment. Input to vertbio.
- detritus(fsm,death\_dia,death\_fla) Computes the phytoplankton death rate as a function of the chlorophyll concentration in the surface layer (see end of Section 3.1.3). The death rates are used as input to biosource.
- oxysat(oxysurf,rho,s,t,fsm) Calculate the concentration of oxygen in the surface layer assuming saturation in contact with one normal atmosphere. Should be called after the transport routine fctf. Used as input to vertbio.
- $phyto_growth(i,j,t,proddia,prodfla,reddia,redfla)$  The main biochemical routine. Computes the phytoplankton growth (3.7), the production maximum (3.10) and the respiration (3.11) at temperature t, for both diatoms and flagellates. The prod and red terms are input to biosource.
- slim(alpha,pmax,s) Computes the modified Michaelis-Menten limitation,  $V_i \in [0...1)$ , for substance S in Equation (3.8). Slim is a local function called by phyto\_growth.

#### 3.3.3 Transport routines

The last seven routines are mainly dealing with the horizontal and vertical distribution (through advection and diffusion) in the water masses of the chemical-biological variables, according to the updates (in time) given by the differential equations in Section 3.2. Based on the old (last time step) concentrations and the new updates of the physical variables, the advection and horizontal diffusion is first updated in **fctf**. Then all source terms from the biochemical routines are included before vertical diffusion is applied in **vertbio**. Finally there are updates due to sinking and the boundary conditions.

- **biobound(icode,ndate,fsm)** Relax the solution of the advection and diffusion equation (and possible sinking, sedimentation and resuspension) in the FRS-zone according to the external specified open boundary condition. Zero the fields in land point.
- deadbot(dia,fla,det,sis) Boundary conditions for algae in the sediment layer. Called after applying the boundary conditions in **biobound**. Tranfer all diatoms and flagellates at the bottom to detritus, and diatoms to also from biogenic silica.

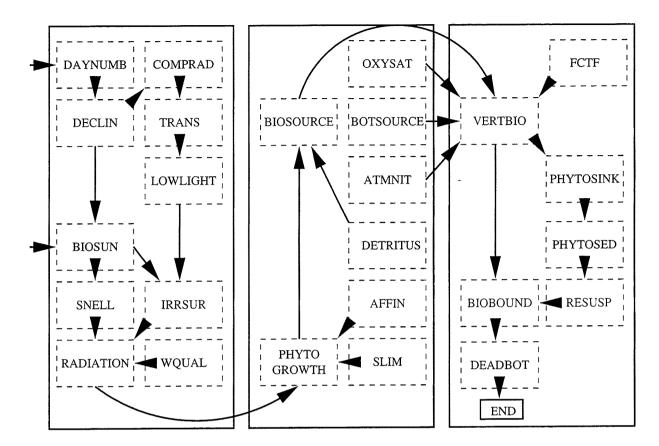


Figure 3.3: Flowchart for the main biology routine, BIOLOGY.F90. Light routines (left), biochemical routines (middle) and transport routines (right)

- fctf(....) Advects the scalar fields, Equation 3.15, and adds the horizontal part of the diffusion, i.e. the second and third term of (3.16), using the flux-corrected transport algorithm. This method is chosen because of the absolute requirement of having positive definite chemical/biological fields. The horizontal diffusivity,  $A_H$ , should be computed by the physical module. After calling fctf, a routine for adding vertical diffusion and for including the surface boundary conditions and source terms must be called, i.e. vertbio.
- phytosink(icode,tmp,d,dz,dzz,dti) Performs the sinking of the particulate matters (DIA, FLA, DET, SIS and ISPM). The sinking of all matters are done after the updates from vertical diffusion in vertbio.
- phytosed(icode,tmp,d,dz,dzz,dti) Performs sedimentation of dead and living algae from the next to bottom layer to the bottom. Called after phytosink.
- resusp(icode,tmp,d,dz,dzz,dti) Performs resuspension of particles from the bottom. Called after phytosed.

**vertbio(....)** Adds the contribution from the vertical diffusion term when advecting scalar fields. The vertical diffusivity,  $K_H$ , should be taken from the physical module. A call to **vertbio** must be preceded by a call to a routine, **fctf**, for advecting horizontally taking into account all terms of the advection equation except the vertical diffusion term. The vertical integration is implicit from time step n to time step n+1. The **source** terms from several biochemical routines are included in the fields before the integration.

#### 3.3.4 Include files

The include-file for the chemical-biological variables is named **biocom.h.** It contains the fields for the seven prognostic variables NIT, PHO, SIL, DET, FLA, DIA and RAD, together with some important constants. See Appendix A for a further description and for an example of this include file.

A second include file, **sedcom.h**, is used for OXY, SIS, ISPM, and all parameters controlling the sedimentation and resuspension. An example of this file can be found in Appendix B.

### 3.4 The Chemical-Biological variables

In this section the complete list of all chemical-biological constants, and variables that goes in and out of the subroutines, are given.

#### GLOBAL PHYSICAL CONSTANTS :

im: Maximum number of points in x-direction

jm: Maximum number of points in y-direction

kb: Maximum number of points in z-direction

#### **REAL CONSTANTS** :

- AA(6): Consists of several constants used for phytoplankton production and respiration. All constants are initiated in AFFIN.F.
  - **aa(1):** Diatom production maximum at  $0 \ C \ (s^{-1})$
  - **aa(2):** Temperature dependent PMAX for diatoms,  $(\mathcal{C}^{-1})$
  - **aa(3):** Flagellate production maximum at  $0 \ C \ (s^{-1})$
  - **aa**(4): Temperature dependent PMAX for flagellates,  $(^{\circ}C^{-1})$ )
  - **aa(5):** Respiration rate at  $0 \mathcal{C}(s^{-1})$
  - **aa(6):** Respiration rate temperature dependence,  $(\mathcal{C}^{-1})$
- **b2:** Extinction due to water and non chlorophyll materials  $(m^{-1})$ . Initiated in BIO-COM.H

CC(4): More phytoplankton constants. Initiated in the include file BIOCOM.H.

- cc(1): Fraction of phosphate and nitrate in a cell (mgP/mgN)
- cc(2): Fraction of silicate and nitrate in a cell (mgSi/mgN)
- cc(3): Phytoplankton death rate,  $(s^{-1})$
- cc(4): Decomposition rate detritus,  $(s^{-1})$
- CCAR: Atomic weight carbon. Initiated in SEDCOM.H

COXY: Atomic weight oxygen. Initiated in SEDCOM.H

COXY2: Atomic weight oxygen gas. Initiated in SEDCOM H

CNIT: Atomic weight nitrate. Initiated in BIOCOM.H

- CPHO: Atomic weight phosphate. Initiated in BIOCOM.H
- CSIL: Atomic weight silicate. Initiated in BIOCOM.H
- **DENITFR:** Fraction of nitrogen in detritus in the sediment that denitrifies. Initiated in SEDCOM.H
- DETBUL: Max amount of DET before burying occurs. Instated in SEDCOM.H
- **DETMAX:** Amount detritus that corresponds to max inorganic nitrogen flux from sediment. Initiated in SEDCOM.H
- **DETW:** Molecule weight DET. Used in connection with TOTSED calculations. Initiated in SEDCOM.H
- **DIAMIN:** Minimum concentration  $(mgN/m^3)$  of diatoms. Initiated in BIOCOM.H. Used in order to prevent the algae in the model to extinct
- **FLAMIN:** Minimum concentration  $(mgN/m^3)$  of flagellates Initiated in BIO-COM.H. Used in order to prevent the algae in the model to extinct.
- **KN\_DIA:** Affinity nitrogen, diatoms  $(\mu MN)^{-1} s^{-1}$ . Initiated in AFFIN.F.

**KN\_FLA:** Affinity nitrogen, flagellates  $(\mu MN)^{-1} s^{-1}$ . Initiated in AFFIN.F.

- **KP\_DIA:** Affinity phosphate, diatoms  $(\mu MP)^{-1} s^{-1}$ . Initiated in AFFIN.F.
- **KP\_FLA:** Affinity phosphate, flagellates  $(\mu MP)^{-1} s^{-1}$ . Initiated in AFFIN.F.
- **KR\_DIA:** Affinity radiation, diatoms  $m^2$  (µEinstein)<sup>-1</sup>. Initiated in AFFIN.F.
- **KR\_FLA:** Affinity radiation, flagellates  $m^2$  (µEinstein)<sup>-1</sup>. Instanted in AFFIN.F.
- **KS\_DIA:** Affinity silicate, diatoms  $(\mu MSi)^{-1} s^{-1}$ . Initiated in AFFIN.F.
- MY: Cosine of the mean zenith angle of the diffuse light. Initiated in BIOCOM.H.
- N2CHLA: Fraction of nitrate and chlorophyll\_a in a cell (mgN/mgCHL\_A). Initiated in BIOCOM.H.
- **NIW:** Molecule weight of N in DET. Used in connection with TOTSED calculations. Initiated in SEDCOM.H

- NY: Chlorophyll light extinction coefficient  $(m \ mgChl_a)^{-1}$ . Initiated in BIO-COM.H.
- **PAR:** Photosynthetic active irradiance (%). Initiated in BIOCOM.H.
- SCC(7): Several constants for oxygen and sedimentation-resuspension. Values set in SEDCOM.H.
  - scc(1): Fraction of oxygen and nitrate for each cell produced (mgO/mgN)
  - scc(2): Fraction of carbon and nitrate in a cell (mgC/mgN)
  - scc(3): Fraction Oxygen/Nitrogen for the denitrification

scc(4): decomposition rate biogenic silica  $(s^{-1})$ 

- scc(5): Minimum rate of leakage of nutrients from sediments  $(s^{-1})$
- scc(6): Maximum rate of leakage of nutrients from sediments  $(s^{-1})$
- scc(7): Burial rate DET and SIS in sediments  $(s^{-1})$
- SIB: Concentration of SIL  $(\mu M)$  when max sinking speed of DIA occurs. Initiated in SEDCOM.H
- SISBUL: Max amount of SIS before burying occurs. Initiated in SEDCOM.H
- SISMAX: Amount biogenic silica that corresponds to max inorganic silicate flux from sediment. Initiated in SEDCOM.H
- SISW: Molecule weight SIS. Used in connection with TOTSED calculations. Initiated in SEDCOM.H
- SIW: Atomic weight silicon. Used in connection with TOTSED calculations. Initiated in SEDCOM.H
- **SRDET:** Sinking rate DET  $(ms^{-1})$ , set in SEDCOM.H
- **SRDIAMAX:** Minimum sinking rate DIA  $(ms^{-1})$ , set in SEDCOM.H

**SRDIAMIN:** Maximum sinking rate DIA  $(ms^{-1})$ , set in SEDCOM.H

**SRFLA:** Sinking rate FLA  $(ms^{-1})$ , set in SEDCOM.H

**SRSED:** Sinking rate ISPM  $(ms^{-1})$ , set in SEDCOM.H

**SRSIS:** Sinking rate SIS  $(ms^{-1})$ , set in SEDCOM.H

**T0:** Temperature for which the half saturation constants are given, see (3.9). Used and initiated in AFFIN.F.

#### LOGICAL VARIABLES :

**BIO:** Set true if biology module should be called

**LOW:** Set true if GLOBAL  $\leq$  MEANGLO

SEDIMENT: Set true if sedimentation should be included

WAVES: Set true if waves should be included

#### **REAL VARIABLES** :

**BEAMTR:** Beam transmittance  $(\mu Einstein/m^2/s)$ 

DAYNR: The day number (1-365, not including February 29)

**DIFFTR:** Diffuse transmittance  $(\mu Einstein/m^2/s)$ 

GLOBAL: Total global irradiance from data

MAXGLO: Global irradiance using clear sky constants

MEANGLO: Global irradiance using mean sky constants

PHI: Zenith angle in water of the direct light

**THETA:** The declination of the sun

**TTIME:** Time of day

#### 2-DIMENSIONAL REAL ARRAYS :

**DEATH\_DIA(IM,JM):** Actual death-rates diatoms  $(s^{-1})$  in each water column. Equals 0 if the concentration of diatoms in the surface cell < DIAMIN

DEATH\_FLA(IM,JM): Same purpose as DEATH\_DIA

DENIT(IM,JM): Accumulated amount of nitrogen denitrified in sediments

DETBU(IM,JM): Accumulated amount of detritus buried in sediments

**DIFFUS(IM,JM):** Diffuse surface irradiance ( $\mu Einstein/m^2/s$ ).

**DIRECT(IM,JM):** Direct surface irradiance ( $\mu Einstein/m^2/s$ ).

**KAPPA(IM,JM):** Coefficient of light extinction depending on the water quality (except for self-shading)

LATIT(IM,JM): Latitude

LONGIT(IM,JM): Longitude

SISBU(IM,JM): Accumulated amount of biogenic silica buried in sediments

SUMDF(IM,JM): Sum (daily) of the diffuse irradiance functions when GLOBAL  $\leq$  MEANGLO

SUNHEI(IM,JM): Sun height

SUNTI(IM,JM): Sun time

**TOTSED**(**IM**,**JM**): Total mass of the accumulated sediment in each gridpoint. Used in the calculation of the resuspension from Equation 3.14.

TTAU(IM,JM): The total bottom stress

#### **3-DIMENSIONAL REAL ARRAYS** :

**DET(IM,JM,KB):** Detritus concentration  $(mgN/m^3)$ 

**DIA**(**IM**,**JM**,**KB**): Diatom concentration  $(mgN/m^3)$ 

**FLA(IM,JM,KB):** Flagellate concentration  $(mgN/m^3)$ 

ISPM(IM,JM,KB): Inorganic suspended particulate matter variable concentration  $(mg/m^3)$ 

**NIT(IM,JM,KB):** Nitrate concentration  $(mgN/m^3)$ 

**OXY(IM,JM,KB):** Oxygen concentration  $(mgO/m^3)$ 

**PHO(IM,JM,KB):** Phosphate concentration  $(mgP/m^3)$ 

**PROD\_DIA**(IM,JM,KB): Production of diatoms  $(mgN/m^3/s)$ 

**PROD\_FLA(IM,JM,KB):** Production of flagellates  $(mgN/m^3/s)$ 

**RAD(IM,JM,KB):** Irradiance in the water column ( $\mu Einstein/m^2/s$ ).

**RED\_DIA(IM,JM,KB):** Respiration of diatoms  $(mgN/m^3/s)$ 

**RED\_FLA(IM,JM,KB):** Respiration of flagellates  $(mgN/m^3/s)$ 

SIL(IM,JM,KB): Silicate concentration  $(mgSi/m^3)$ 

SIS(IM,JM,KB): Biogenic silica concentration  $(mgSi/m^3)$ 

SOURCE(IM,JM,KB): Source terms for an actual chemical or biological constituent  $(mg/m^3)$ 

### **3.5** User supplied input/initialization

The user must support initial fields for the prognostic variables NIT, PHO, SIL, DET, FLA, DIA, OXY, ISPM and SIS, and fields for the inflow of these variables on the open boundaries should also be supplied. These initial and boundary conditions can either be given in separate user supplied routines, or together with the initial and boundary conditions for the physical part of the model. The user must also supply nutrient input from rivers and the atmosphere together with the river fluxes and temperature.

The chemical-biological model depends on a number of constants, and the output from the model will reflect the way these are initialized. The constants are set in the routines **biocom**, sedcom and affin. The routine fctf uses values for horizontal diffusivities, and vertbio a vertical diffusion coefficient. These values are related to similar physical constants, and should be supplied by the physical model. The biological module also needs fields for bottom stress to be used in the sedimentation and resuspension routines. For these computations wave fields should be included.

Vertbio needs a field for surface fluxes or values. In the model these are all set to zero, except for the oxygen concentration, where saturation in contact with one normal atmosphere is assumed.

The light routines could be substituted by real light if that is available. Several constants (see Eqs. (3.2) and (3.3), and subroutines are used for the computations (Skartveit & Olseth, 1986, 1987). These can be varied to give a more sunny/cloudy weather, dependent of the geographical area to be modeled. The constants set in the model at present are for Bergen(3.2), and the average of Bergen and Montreal (3.3).

Parameter	Explanation	Value
a(1)	diatom production maximum at $0  \mathcal{C}$	$1.53\text{E-5}~(s^{-1})$
a(2)	diatom temperature dependent $P_{max}$	$0.063 \ (^{\circ}C^{-1})$
a(3)	flagellate production maximum at $0{ m C}$	$1.02\text{E-5}~(s^{-1})$
a(4)	flagellate temperature dependent $P_{max}$	$0.063 \ ({}^{\circ}\!C^{-1})$
a(5)	metabolic loss rate at $0{}^{\circ}\!C$	$8.05\text{E-7}~(s^{-1})$
a(6)	metabolic loss rate temp. dependence	$0.07 \ (C^{-1})$
$lpha_{d.1}$	growth affinity for irradiance (diatoms)	$3.6\text{E-7}\ m^2\ \mu Einstein^{-1}$
$lpha_{d.2}$	growth affinity for nitrate (diatoms)	$1.7\text{E-5}\ s^{-1}\ \mu M^{-1}$
$\alpha_{d.3}$	growth affinity for phosphate (diatoms)	$2.7\text{E-4}\ s^{-1}\ \mu M^{-1}$
$\alpha_{d.4}$	growth affinity for silicate (diatoms)	$2.5\text{E-5}\ s^{-1}\ \mu M^{-1}$
$lpha_{d.1}$	growth affinity for irradiance (flagellates)	$1.1\text{E-7}\ m^2\ \mu Einstein^{-1}$
$lpha_{d.2}$	growth affinity for nitrate (flagellates)	$1.5\text{E-5}~s^{-1}~\mu M^{-1}$
$lpha_{d.3}$	growth affinity for phosphate (flagellates)	$2.5\text{E-}4~s^{-1}~\mu M^{-1}$
b2	Extinction due to water and non chlorophyll	$0.07 \ (m^{-1})$
cc(1)	intercellular $P/N$ relationship	$0.138~(mgPmgN^{-1})$
cc(2)	intercellular $Si/N$ relationship	$1.75 \ (mgSimgN^{-1})$
cc(3)	death rate $(0.1)$	$1.6\text{E-6}~(s^{-1})$
cc(4)	rate of decomposition detritus $(0.013)$	$1.52\text{E-7}~(s^{-1})$
DENITFR	fraction of nitrogen in sediment that denitrifies	0.3
DETBUL	limit of burying of DET	$630 \ (mg \ m^{-3})$
DETMAX	corresp. conc. to max flux of NIT from sediment	$630 \ (mg \ m^{-3})$
DETW	molecule weight DET	3555
DIAMIN	Minimum value DIA concentration	$0.1 \ (mgNm^{-3})$
FLAMIN	Minimum value FLA concentration	$0.1 \ (mgNm^{-3})$
$\mu$	mean cosine of diffuse light zenith angle	0.83
N2CHLA	cellular fraction of nitrate and Chl_a	$11.0 \ (mgN \ mgChl\_a^{-1})$
NIW	molecule weight of N in DET	224
ν	Chl_a light extinction coefficient	$1.38\text{E-}2 \ (m  mg Chl_a^{-1})$
PAR	photosynthetic active irradiance	40 %

.....continues on next page .....

SISBUL	limit of burying of SIS	$6720 \ (mgm^{-2})$
SISMAX	corresp. conc. to max flux of SIL from sediment	$6720 \ (mgm^{-2})$
SISW	molecule weight SIS	60
SIW	atomic weight silicon	28
SRDET	Sinking rate DET	$3.0 \ (md^{-1})$
SRDIA <sub>min</sub>	Minimum sinking rate DIA	$0.3 \ (md^{-1})$
SRDIA <sub>max</sub>	Maximum sinking rate DIA	$3.0 \ (md^{-1})$
SRFLA	Sinking rate FLA	$0.25 \ (md^{-1})$
SRSED	Sinking rate ISPM	$1.0 \ (md^{-1})$
SRSIS	Sinking rate SIS	$3.0 \ (md^{-1})$
SIB	Conc. of SIL when max sinking speed DIA	$1.0~(\mu M)$
$\operatorname{scc}(1)$	Fraction of OXY and NIT for each cell produced	19.71~(mgO/mgN)
$\operatorname{scc}(2)$	Fraction of C and NIT in a cell	5.68~(mgC/mgN)
$\operatorname{scc}(3)$	Fraction OXY/NIT for the denitrification	3.42
$\operatorname{scc}(4)$	Decomposition rate biogenic silica	$1.45\text{E-8}~(s^{-1})$
$\operatorname{scc}(5)$	Min. leakage rate of nutrients from sediments	$1.157\text{E-6}~(s^{-1})$
$\operatorname{scc}(6)$	Max. leakage rate of nutrients from sediments	$4.63\text{E-}6~(s^{-1})$
$\operatorname{scc}(7)$	Burial rate (120 days)	9.65E-8 $(s^{-1})$

Table 3.1: ...(continue)... Parameter values

# Chapter 4

## Some model results and examples

In this chapter a few example figures are given, to show some of the application that the model has been used for. The results are not properly discussed, for that purpose the papers and reports referred should be consulted.

For most purposes NORWECOM has been used in an area covering an extended North Sea with a horizontal resolution of  $20 \times 20$  kilometers. This resolution is too coarse to proper model processes within the Skagerrak, and for that purposes a nested version of the model is used. In the nested model, boundary values from the coarse North Sea model is used as input to a fine grid ( $4 \times 4$  kilometer) model for the Skagerrak and Kattegat. These boundaries are updated every hour, to resolve tides. In both models 12  $\sigma$ -layers have been used, with a relative finer resolution in the surface layers to better resolve the biological and chemical processes that are most important here. The two bottom topographies are shown in Figure 4.1.

Using the physical module of NORWECOM the North Sea circulation, and it's long term variability, has been modeled (e.g. Smith *et al.* (1996); Iversen *et al.* (1998)). In Figure 4.2 the modeled mean (1976-1996) winter circulation at 10 meters depth in the North Sea is shown, while in Figure 4.3 the variability in the modeled inflow through a section from the Orkneys to Norway (along 59.17 °N) is given.

In Skogen *et al.* (1995) the model was for the first time used to model a whole annual cycle of the North Sea primary production. Results for individual ICES boxes were reported, and compared with measurements. In addition a scenario where the anthrophogenic inputs of inorganic nitrogen and phosphorus were reduced with 50 %, to measure the effect of such reduction in accordance with agreements from the International conference on the protection of the North Sea (London 1987). The results, annual integrated primary production  $(gC/m^2/year)$  and percentage reduction are given in Figure 4.4.

To study the dispersion and dilution of different water masses, it is possible to label these in the model. This is done by including an extra prognostic variable, in the model, and treat it as a passive tracer by the modeled circulation and diffusion. This algorithm was used in Skogen *et al.* (1997b), where the transport of water with origin in the German Bight were labeled, to investigate this transport to Skagerrak. In this work the nested model system (North Sea  $\rightarrow$  Skagerrak) was used, and the model was run several years to

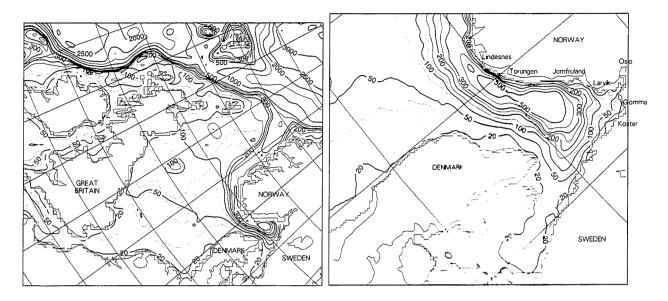
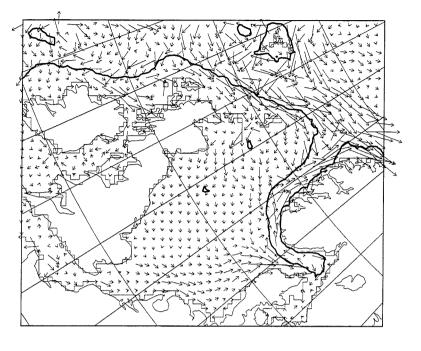


Figure 4.1: Bottom topography, North Sea  $20 \times 20$  km. left, and Skagerrak  $4 \times 4$  km. right

look into variability between different years. In Figure 4.5 the interannual variability of water from the German Bight through a section (Tyboron) on the Damsh west coast are shown.

Another eutrophication issue is the concern about oxygen depletion. In Skogen (1998b) different nutrient reduction strategies was investigated in the light of change in minimum bottom oxygen values in the German Bight. The change in this minimum was discussed, as freshwater nutrients were reduced from either the Rhine or the German rivers (Elbe, Ems and Weser) - or both. In Figure 4.6 oxygen in the bottom layer for a reference run is given.

One validation process that the model has been through, is a comparison with climatological nutrient fields in the North Sea (Søiland & Skogen, 1998). Using a costfunction (Berntsen *et al.*, 1996), where the absolute difference between model and data are normalized using the standard deviation, quantitative measures on the model performance can be established. In Figure 4.7 such an example is given. Both the modeled phosphate field, and the costfunction for this field is shown.



→ (10 cm/s)

Figure 4.2: Modeled mean (1976-1996) flow field for January, February and March at 10 meters depth. The solid line is the 200 meters depth contour

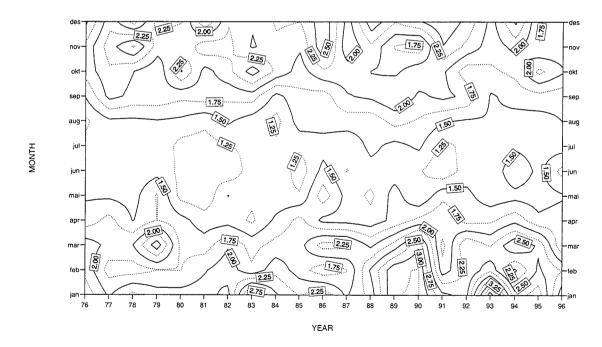


Figure 4.3: Modeled transports (in Sverdrup) southwards through section Orkney - Norway (Utsira) for the period 1976-1996

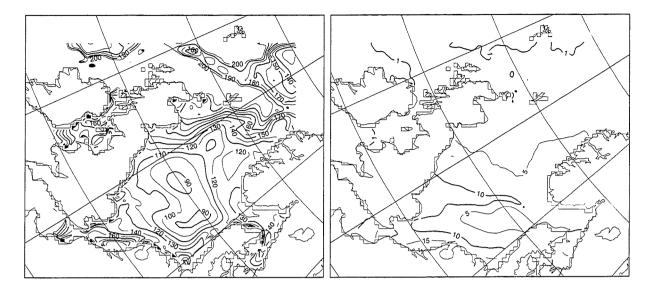


Figure 4.4: Annual depth integrated primary production  $(gC/m^2/year)$  for 1985 (left), and reduction (in percentage) with a 50 % reduction of anthrophogenic inputs (right)

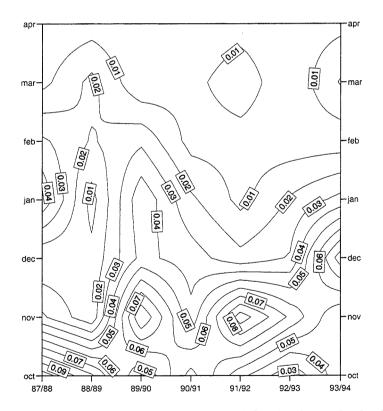


Figure 4.5: Monthly mean transport in Sverdrup of GBW through the Tyborøn section

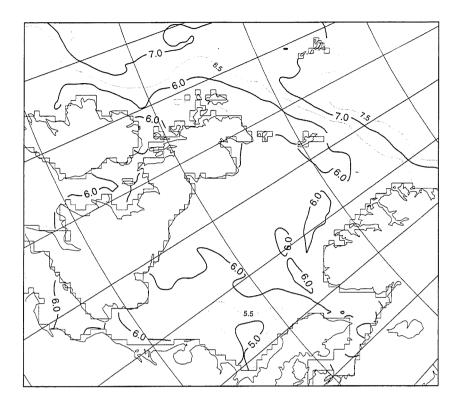


Figure 4.6: Oxygen (ml/l) in the bottom layer (2.5 % of total depth above bottom) at July 1 in the reference run

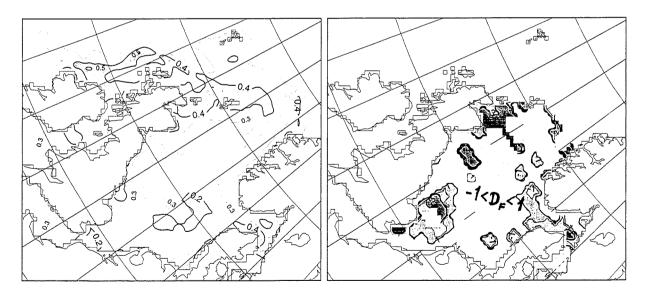


Figure 4.7: Mean (1980-89) model (left) and costfunction (right) field for phosphate. The results are for May-June-July in the upper 20 meters. The nutrient fields are in  $(\mu M)$ , while the isolines in the costfunction field are -3, -2, -1, +1, +2 and +3

.-

## Appendix A The include file BIOCOM

This appendix gives an example of an include file, BIOCOM.H, used together with the chemical-biological part of NORWECOM. The include file consists of common blocks defining the seven prognostic variables (NIT, PHO, SIL, DET, FLA, DIA and RAD) together with some important constants.

```
\mathbf{C}
C Atomic weights
\mathbf{C}
\mathbf{C}
     CNIT = nitrate
\mathbf{C}
     CPHO = phosphate
\mathbf{C}
     CSIL = silicate
\mathbf{C}
       REAL CNIT, CPHO, CSIL
       DATA CNIT/14.01/,CPHO/30.97/,CSIL/28.09/
\mathbf{C}
\mathbf{C}
     CC(1): Fraction of phosphate and nitrate in a cell (mgP/mgN)
\mathbf{C}
     CC(2): Fraction of silicate and nitrate in a cell (mgSi/mgN)
\mathbf{C}
     CC(3): Phytoplankton death rate (s^{-1})
     CC(4): Decomposition rate detritus (s^{-1})
\mathbf{C}
\mathbf{C}
       REAL CC(4)
       DATA CC(1)/0.138/,CC(2)/1.75/,CC(3)/1.6E-6/,CC(4)/1.52E-7/
\mathbf{C}
C Minimum values of phytoplankton concentrations (mgN/m^3)
\mathbf{C}
       REAL DIAMIN, FLAMIN
       PARAMETER(DIAMIN=0.1,FLAMIN=0.1)
\mathbf{C}
     N2CHLA : Fraction of nitrate and Chl_A in a cell. (mgN/mgChl_A)
\mathbf{C}
\mathbf{C}
     MY : Cosine of the mean zenith angle of the diffuse light.
     NY : Chlorophyll light extinction coefficient (m \ mqChl_A)^{-1}.
\mathbf{C}
```

C PAR : Photosynthetic active radiance (%)

## APPENDIX A. THE INCLUDE FILE BIOCOM

**B2**: Extinction due to water  $(m^{-1})$  $\mathbf{C}$ RC1 : Refraction coefficient air С  $\mathbf{C}$ RC1 : Refraction coefficient water  $\mathbf{C}$ REAL MY,NY,PAR,N2CHLA,B2,RC1,RC2 PARAMETER (MY=0.83,NY=1.38E-2,PAR=0.40,N2CHLA=11.0, +B2=0.07, RC1=1.0, RC2=1.34) $\mathbf{C}$ C Nutrients common declarations  $\mathbf{C}$ REAL NIT(IM,JM,KB),PHO(IM,JM,KB),SIL(IM,JM,KB) COMMON /NUTCOM/NIT,PHO,SIL  $\mathbf{C}$ C Biological common declarations  $\mathbf{C}$ REAL DET(IM,JM,KB),FLA(IM,JM,KB),DIA(IM,JM,KB) COMMON /BIOCOM/DET,FLA,DIA  $\mathbf{C}$ C Light common declaration  $\mathbf{C}$ REAL RAD(KB) COMMON /LIGHT/RAD

34

## Appendix B The include file SEDCOM

This appendix gives an example of an include file, SEDCOM.H, used together with the chemical-biological part of NORWECOM. The include file consists of common blocks defining the three new prognostic variables from version 1.0 (SIS, OXY, ISPM), together with constants used in the sedimentation-resuspension routines.

```
\mathbf{C}
C Sediment common declarations
\mathbf{C}
      REAL TTAU(IM,JM)
      LOGICAL WAVES, SEDIMENT
      COMMON/COMSED/TTAU
      COMMON/COMLOG/WAVES,SEDIMENT
\mathbf{C}
      REAL SIS(IM,JM,KB),OXY(IM,JM,KB),ISPM(IM,JM,KB)
      COMMON /BIOSED/SIS,OXY,ISPM
\mathbf{C}
      REAL TOTSED(IM,JM)
      REAL DETBU(IM,JM),DENIT(IM,JM),SISBU(IM,JM)
      COMMON/BOTW/TOTSED
      COMMON/BOTPRO/DETBU, DENIT, SISBU
\mathbf{C}
C Atomic weights and molecule weights
\mathbf{C}
      REAL DETW, SISW, NIW, SIW
      PARAMETER (DETW=3555., SISW=60., NIW=224., SIW=28.)
\mathbf{C}
\mathbf{C}
    COXY = oxygen
\mathbf{C}
    COXY2 = oxygen gas (O_2)
\mathbf{C}
    CCAR = carbon
\mathbf{C}
      REAL COXY,COXY2,CCAR
      DATA COXY/16.00/,COXY2/32.00/,CCAR/12.01/
```

 $\mathbf{C}$ 

- C SCC(1): Fraction of oxygen and nitrate for each cell produced (mgO/mgN)
- C SCC(2) : Fraction of carbon and nitrate in a cell (mgC/mgN)
- C SCC(3): Fraction Oxygen/Nitrogen for the denitrification
- C SCC(4) : Decomposition rate biogenic silica-180 days  $(s^{-1})$
- C SCC(5): Minimum rate of leakage of nutrients from sediments 10 days  $(s^{-1})$
- C SCC(6): Maximum rate of leakage of nutrients from sediments 6 hours  $(s^{-1})$
- C SCC(7) : Burial rate 120 days  $(s^{-1})$

 $\mathbf{C}$ 

REAL SCC(6)

DATA SCC(1)/19.71/,SCC(2)/5.68/,SCC(3)/3.42/,SCC(4)/1.45E-8/ DATA SCC(5)/1.157E-6/,SCC(6)/4.63E-5/,SCC(7)/9.65E-8/ REAL DETMAX,DETBUL,DENITFR,SISBUL,SISMAX DATA DENITFR/0.3/,DETMAX/630./,DETBUL/630./ DATA SISMAX/6720./,SISBUL/6720./

 $\mathbf{C}$ 

C Sinking speeds

 $\mathbf{C}$ 

INTEGER DAY

```
REAL SRDET,SRFLA,SRDIAMAX,SRDIAMIN,SRSIS,SIB,SRSED
PARAMETER(DAY=86400,SIB=1.0,SRDET=3.,SRFLA=0.25)
PARAMETER(SRSED=1.0,SRDIAMAX=3.0,SRDIAMIN=0.3,SRSIS=3.0)
```

## Bibliography

- AKSNES, DAG L., & LIE, ULF. 1990. A coupled Physical-Biological Pelagic model of a shallow Sill Fjord. *Estuarine, Coastal and Shelf Science*, **31**, 459–486.
- AKSNES, D.L., EGGE, J.K., ROSLAND, R., & HEIMDAL, B.R. 1994. On the representation of Emilania huxleyi in phytoplankton simulation models - A first approach. Sarsia, 79, 291–300.
- AKSNES, D.L., ULVESTAD, K.B., BALIÑO, B., BERNTSEN, J., EGGE, J., & SVENDSEN,
  E. 1995. Ecological modelling in coastal waters : Towards predictive Physical-Chemical-Biological simulation models. *Ophelia*, 41, 5–36.
- ANDERSEN, VALERIE, & NIVAL, PAUL. 1989. Modelling phytoplankton population dynamics in an enclosed water column. Journal of the Marine Biological Association of the UK, **69**, 625-646.
- BERNTSEN, JARLE, SVENDSEN, EINAR, & OSTROWSKI, MAREK. 1996. Validation and Sensitivity study of a Sigma-coordinate Ocean Model using the SKAGEX dataset. ICES C.M.1996/C:5.
- BLUMBERG, A. F., & MELLOR, G. L. 1980. A coastal Ocean Model. Pages 203-214 of: SUNDERMAN, J., & HOLTZ, K.P. (eds), Mathematical Modelling of Estuarine Physics. Springer-Verlag, Berlin.
- BLUMBERG, A.F., & MELLOR, G.L. 1987. A description of a three-dimensional coastal ocean circulation model. *In:* HEAPS, N. (ed), *Three-Dimensional Coastal Ocean Models*, *Vol.4*. American Geophysical Union.
- Cox, M.D., & BRYAN, K. 1984. A Numerical Model of the Ventilated Thermocline. J. of Physical Oceanography, 14, 674–687.
- DANIELSSEN, D. S., EDLER, L., FONSELIUS, S., HERNROTH, L., OSTROWSKI, M., SVENDSEN, E., & TALPSEPP, L. 1997. Oceanographic variability in Skagerrak/Northern Kattegat, May-June 1990. ICES Journal of Marine Science, 54, 753-773.
- DANIELSSEN, D.S., DAVIDSSON, L., EDLER, L., FOGELQVIST, E., FONSELIUS, S., FØYN, L., HERNROTH, L., HÅKANSON, B., OLSSON, I., & SVENDSEN, E. 1991. SKAGEX: Some preliminary results. 33pp, ICES C.M.1991/C:2.

- DEE, DICK P. 1994. A pragmatic approach to model validation. Pages 1-14 of: LYNCH, D.R., & DAVIES, A.M. (eds), Quantitative Skill Assessment for Coastal Ocean Models. American Geophysical Union. ISBN 0-87590-261-8.
- EIDE, L.I., REISTAD, M., & GUDDAL, J. 1985. Database av beregnede vind og bølgeparametre for Nordsjøen, Norskehavet og Barentshavet, hver 6.time for årene 1955-81. DNMI, Oslo/Bergen, Norway. 38pp, In Norwegian.
- EPPLEY, R. W. 1972. Temperature and Phytoplankton Growth in the Sea. Fishery Bulletin, 70(4).
- GEHLEN, M., MALSCHAERT, H., & RAAPHORST, W.R. 1995. Spatial and temporal variability of benthic silica fluxes in the southeastern North Sea. *Cont. Shelf Res.*, 13, 1675–1696.
- GUSTAFSSON, BO, & JÖNSSON, ANETTE. 1995. Verification of hydrodynamic models applied to Kattegat and Skagerrak. 142pp.
- IVERSEN, S.A., SKOGEN, M.D., & SVENDSEN, E. 1998. Influx of Atlantic water and feeding migration of Horse Mackerel. ICES C.M.1998/R:18.
- LAANE, R.W.P.M., SVENDSEN, E., RADACH, G., GROENEVELD, G., DAMM, P., PÄTSCH, J., DANIELSSEN, D., FØYN, L., SKOGEN, M., OSTROWSKI, M., & KRAMER, K.J.M. 1996. Variability in fluxes of nutrients (N,P,Si) into the North Sea from Atlantic Ocean and Skagerrak caused by variability in water flow. *Deutsche Hydrographische Zeitschrift*, 48(3/4), 401-419.
- LOHSE, L., MALSCHAERT, F.P., SLOMP, C.P., HELDER, W., & RAAPHORST, W. 1995. Sediment-water fluxes of inorganic nitrogen compounds along the transport route of organic matter in the North Sea. *Ophelia*, **41**, 173–197.
- LOHSE, L., KLOOSTECHUIS, H.T., RAAPHORST, W., & HELDER, W. 1996. Denitrification rates as measured by the isotope pairing method and by the acetylene inhibilition technique in continental shelf sediments of the North Sea. *Mar.Eco.Prog.Ser.*, **132**, 169–179.
- MARTINSEN, E. A., ENGEDAHL, H., OTTERSEN, G., ÅDLANDSVIK, B., LOENG, H., & BALIÑO, B. 1992. *MetOcean MOdeling Project, Climatological and hydrographical data for hindcast of ocean currents.* Tech. rept. 100. The Norwegian Meteorological Institute, Oslo, Norway. 93pp.
- MARTINSEN, E.A., & ENGEDAHL, H. 1987. Implementation and testing of a lateral boundary scheme as an open boundary condition in a barotropic ocean model. *Coastal Engineering*, **11**, 603–627.

- MARTINSEN, E.A., SLØRDAL, L.H., & ENGEDAHL, H. 1990. MetOcean MOdeling Project (MOMOP), Phase 2, Data report: Joint presentation of the test cases. Tech. rept. 87. The Norwegian Meteorological Institute.
- MAYER, B. 1995. Ein dreidimensionales, numerisches Scwebstoff-Transportmodell mit Anwendung auf die Deutsche Bucht. Tech. rept. GKSS 95/E/59. GKSS-Forschungszentrum Geesthacht GmbH.
- MELLOR, GEORGE L. 1996 (June). User's guide for a 3-dimensional. primitive equation, numerical ocean model. Prog. in Atmos. and Ocean. Sci, Princeton University, 38pp.
- MELLOR, G.L., & YAMADA, T. 1982. Development of a turbulence closure model for geophysical fluid problems. *Rev. Geophys. Space Phys.*, **20**, 851-875.
- MESINGER, F., & ARAKAWA, A. 1976. Numerical methods used in atmospheric models. Tech. rept. 17. GARP Publication Series.
- O'CONNOR, WILLIAM P. 1991 (March). A User's Manual for the Princeton Numerical Ocean Model. Tech. rept. Institute of Naval Oceanography. Stemms Space Center, MS 39529-5005.
- OEY, L.-Y., & CHEN, P. 1992. A model simulation of circulation in the Northeast Atlantic Shelves and Seas. Journal of Geophysical research, 97(C12), 20087-20115.
- OSPAR. 1998. Report of the ASMO modelling Workshop on Eutrophication Issues 5-8 November 1996, The Hague, The Netherlands.
- OSTROWSKI, MAREK. 1994. The SKAGEX Atlas. A User's Guide Tech. rept. TemaNord 1994:635. Nordic Council of Ministers. Part II : Pp.33-99.
- PLATT, TREVOR, & JASSBY, ALAN D. 1976. The relationship between photosynthesis and light for natural assemblage of coastal marine phytoplankton. J of Phycology, 12, 421-430.
- POHLMANN, T., & PULS, W. 1994. Currents and transport in water Pages 345-402 of: SÜNDERMANN, J. (ed), Circulation and contaminant fluxes in the North Sea. Berlin: Springer Verlag.
- PROCTOR, R., BAART, A., BERG, P., BOON, J., DELEERSNIJDER. E., DELHEZ, E., GARREAU, P., GERRITSEN, H., JONES, J.E., DE KOK, J., LAZURE, P., LUYTEN, P., OZER, J., POHLMANN, T., RUDDICK, K., SALDEN, R., SALOMON, J.C., SKO-GEN, M., TARTINVILLE, B., & VESTED, H.J. 1997. Final report NOMADS NOrth sea Model Advection Dispersion Study. Tech. rept. POL Internal 108. Proudman Oceano-graphic Laboratory. EC concerted action : MAS2-CT94-0105.
- REISTAD, M., EIDE, L.I., GUDDAL, J., & MAGNUSSON, A.K. 1988. Wave model sensitivity study. The Norwegian Meteorolgical Institute.

- RØED, LARS PETTER, HACKETT, BRUCE, & SKÅTUN, H. 1989. Metocean modeling project (MOMOP): Final report: Results of the model intercomparison (a bridged version). Tech. rept. 89-3312. VERITEC.
- SÆTRE, R., SVENDSEN, E., FOSSUM, P., SKOGEN, M.D., & ERIKSRØD, G. 1998. On the site-specific role of the central Norwegian shelf for the recruitment strategy of the Norwegian spring spawning herring. ICES C.M. 1998/R:17.
- SKARTVEIT, A., & OLSETH, J. A. 1986. Modelling slope irradiance at high lattitudes. Solar Energy, 36(4), 333-344.
- SKARTVEIT, A., & OLSETH, J. A. 1987. A model for the diffuse fraction of hourly global radiation. *Solar Energy*, **37**, 271–274.
- SKOGEN, M. D. 1993. A User's guide to NORWECOM, the NORWegian ECOlogical Model system. Tech. rept. 6. Institute of Marine Research, Division of Marine Environment, Pb.1870, N-5024 Bergen. 23pp.
- SKOGEN, M. D., SVENDSEN, E., BERNTSEN, J., AKSNES, D., & ULVESTAD, K.B. 1995. Modelling the primary production in the North Sea using a coupled 3 dimensional Physical Chemical Biological Ocean model. *Estuarine, Coastal and Shelf Science*, 41, 545-565.
- SKOGEN, MORTEN D. 1998a. A biophysical model for the Benguela upwelling system. Accepted in S.Afr.J.of Mar.Sci.
- SKOGEN, MORTEN D. 1998b. Optimizing nutrient reduction strategies in the marine environment - A simplified example from the North Sea. Tech. rept. Fisken og Havet 2/98. Institute of Marine Research, Pb.1870, N-5024 Bergen.
- SKOGEN, MORTEN D., & MOLL, ANDREAS. 1998. Natural variability of the North Sea primary production. Submitted to Cont.Shelf Res.
- SKOGEN, MORTEN D., MONSTAD, TERJE, & SVENDSEN, EINAR. 1997a. A possible separation between a nothern and a southern stock of the North East Atlantic Blue Whiting. Accepted in Fisheries Research.
- SKOGEN, MORTEN D., ERIKSRØD, GRO, & SVENDSEN, EINAR. 1997b. Quantification of transports to Skagerrak. A modelling approach. Pages 327-339 of: ÖZSOY, E., & MIKAELYAN, A. (eds), Sensitivity to change : Black Sea, Baltic Sea and North Sea. Kluwer Academic Publishers.
- SKOGEN, MORTEN D., SVENDSEN, EINAR, & OSTROWSKI, MAREK. 1997c. Quantifying Volume Transports during SKAGEX with the Norwegian Ecological Model system. Cont. Shelf Res., 17(15), 1817–1837.

- SKOGEN, MORTEN D., AURE, JAN, DANIELSSEN, DIDRIK, & SVENDSEN, EINAR. 1998. Natural fertilization of the marine environment. Modeling of the Glomma flood 1995. Accepted in Sarsia.
- SLØRDAL, L.H., MARTINSEN, E.A., & ENGEDAHL, H. 1991. MetOcean MOdeling Project (MOMOP), Phase 2, Final report: Sensitivity tests and pre-operational simulations. Tech. rept. Tech. Rep 88. The Norwegian Meteorological Institute.
- SMAGORINSKY, J. 1963. General circulation experiments with the primitive equations, I. The basic experiment. Mon. Weather Rev., 91, 99–164.
- SMITH, J., DAMM, P.E., SKOGEN, M.D., FLATHER, R.A., & PÄTSCH, J. 1996. An investigation into the Variability of Circulation and Transoprt of the North-West European shelf using three hydrodynamic models. *Deutsche Hydrographische Zeitschrift*, 48(3/4), 325-348.
- SØILAND, H., & SKOGEN, M.D. 1998. Validation of a 3-D biophysical model using nutrient observations in the North Sea. ICES C.M. 1998/Q:3.
- SØILAND, H., SVENDSEN, E., SKOGEN, M.D., & ERIKSRØD, G. 1996. Numerisk modellering av primærproduksjon og transport av vannmasser og næringssalter langs Norskekysten. Tech. rept. Fisken og Havet, 25. Havforskningsinstituttet, Pb.1870, N-5024 Bergen, Norway.
- STIGEBRANDT, A. 1980. Barotropic and baroclinic response of a semi-enclosed basin to barotropic forcing of the sea. Pages 141-164 of: FREELAND, H.J., FARMER, D.M., & LEVINGS, C.D. (eds), Proceeding of the NATO Conference on Fjord Oceanography. Plenum Press, New York.
- SVENDSEN, E., ERIKSRØD, G., & SKOGEN, M.D. 1995a. Kvantifisering av langtransporterte vannmasser fra Tyskebukta, Østersjøen og Nordsjøen til ytre Oslofjord. Tech. rept. Fisken og Havet, 15. Havforskningsinstituttet, Pb.1870, N-5024 Bergen, Norway.
- SVENDSEN, E., ERIKSRØD, G., & SKOGEN, M.D. 1995b. Numerisk modellering av transport av næringssalter og primærproduksjon i Skagerrak/Kattegat og Ytre Oslofjord. Tech. rept. Fisken og Havet, 28. Havforskningsinstituttet, Pb.1870, N-5024 Bergen, Norway.
- SVENDSEN, E., FOSSUM, P., SKOGEN, M.D., ERIKSRØD, G., BJØRKE, H., NEDRAAS, K., & JOHANNESSEN, A. 1995c. Variability of the drift patterns of Spring Spawned herring larvae and the transport of water along the Norwegian shelf. ICES C.M. 1995/Q:25.
- SVENDSEN, E., BERNTSEN, J., SKOGEN, M.D., ÅDLANDSVIK, B., & MARTINSEN, E. 1996a. Model simulation of the Skagerrak circulation and hydrography during SKAGEX. J. of Mar. Syst., 8(3-4), 219-236.

SVENDSEN, E., SKOGEN, M.D., MONSTAD, T., & COOMBS, S. 1996b. Modelling the variability of the drift of blue whiting larvae and its possible importance for recruitement. ICES C.M. 1996/S:31.

SWAMP-GROUP. 1985. Ocean wave modelling. Plenum Press.

WALSH, M., SKOGEN, M.D., REID, D.G., SVENDSEN, E., & MCMILLAN, J.A. 1996. The relationship between the location of Western Mackerel spawning. larval drift and recruit distributions : A modelling study. ICES C.M. 1996/S:33.