

## Reply to comments by topical editor:

*Comments to the Author:*

*Thank you for your response to your referees. I have reviewed this and have a few suggestions concerning the proposed modifications to the manuscript.*

*A general remark I would make is that there are several places in your response where you cite either unpublished or grey literature material in support of your changes to the manuscript. I would instead suggest that this manuscript is an ideal venue for such issues to be properly raised, both to support this manuscript and to provide a formally published source for these issues that would be useful for future iterations of this model.*

Reply: You are right, we are using a lot of grey litterateur. We have located the ‘grey’ literature that we cite and evaluated in each case if it is possible to use peer-review literature instead. It seems though, that especially for the datasets and models from before journals where available where one could publish such descriptions it is hard to avoid citing reports. We also made sure that this literature is available online.

Carnes, M. R.: Description and Evaluation of GDEM-V 3.0, 2009.

- In this case we could not find an peer- review publication for this version of the GDEM-clintology.

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Conkright, M. E., Antonov, J. I., Baranova, O., Boyer, T. P., Garcia, H. E., Gelfeld, R., Johnson, D. D., Locarnini, R. A., Murphy, P. P., O’Brien, T. D., Smolyar, I. and Stephens, C.: World Ocean Database 2001, Volume 1: Introduction, U.S. Government Printing Office, Washington, D.C., 2002.

- We were not successfull in finding a peer review publcation for this dataset, but we had included the wrong reference, it should be “Conkright, M. E., Locarnini, R. a, Garcia, H. E., O’Brien, T. D., Boyer, T. P., Stepens, C. and Antonov, J. I.: World Ocean Atlas 2001: Objective analyses, data statistics, and figures CD-ROM documentation, Natl. Oceanogr. Data Cent. Intern. Rep. (NOAA Atlas NESDIS), 17(September), 17, 2002“. This has been ammended.

Hansen, C.: Simulated primary production in the Norwegian Sea - Interannual variability and impact of mesoscale activity, Univeristy of Bergen., 2008.

- See answer to specific comment below.

Jaccard, P., Ledang, A., Hjermann, D., Reggiani, E., Sørensen, K., Wehde, H., Kaitala, S. and Folkestad, A.: Real Time Quality Control of biogeochemical measurements, MyOcean document., 2014.

- This reference is not easily accessible, so we have chosen to remove the entire sentence.

Ottersen, G., Aasen, S. and Aure, J.: Utarbeidelse av et program for overvåkning av et eutrofitilstand og –utvikling i norske kystfarvann basert på både tradisjonelle og høyteknologiske metoder, Fisk. og Havet, (1), 144, 1998.

-This reference is easily accessible online, and has never been published in english, so we would prefer to keep it like it is. We have removed the 'e.g' in front and now also refer to a specific figure in the report.

Pätsch, J., Kühn, W., Moll, A. and Lenhart, H.: ECOHAM4 User Guide - Ecosystem Model, Hamburg, Version 4. Technical Reports 01-2009, Institut für Meereskunde, Hamburg, Germany., 2009.

- This reference was changed to: Pätsch, J. and Kühn, W.: Nitrogen and carbon cycling in the North Sea and exchange with the North Atlantic - a model study, Part I. Nitrogen budget and fluxes, Cont. Shelf Res., 28(6), 767–787, 2008.

Samuelsen, A. and Bertino, L.: Arctic Ocean ecosystem modeling in MyOcean, Mercat. Ocean Q. Newsl., 40, 36–44, 2011.

- This is the only publication where we apply this version of the model, so we keep this reference.

Simmons, A. S., Uppala, D. D. and Kobayashi, S.: ERA-Interim: New ECMWF reanalysis products from 1989 onwards, 2007.

- This reference has been changed to: Dee, D. P., Uppala, S. M., Simmons, A. J., Berrisford, P., Poli, P., Kobayashi, S., Andrae, U., Balmaseda, M. A., Balsamo, G., Bauer, P., Bechtold, P., Beljaars, A. C. M., Van De Berg, L., Bidlot, J., Bormann, N., Delsol, C., Dragani, R., Fuentes, M., Geer, A. J., Haimberger, L., Healy, S. B., Hersbach, H., Hólm, E. V, Isaksen, I., Kållberg, P., Köhler, M., Matricardi, M., McNally, A. P., Monge-Sanz, B. M., Morcrette, J. J., Park, B. K., Peubey, C., de Rosnay, P., Tavolato, C., Thépaut, J. N. and Vitart, F.: The ERA-Interim reanalysis: configuration and performance of the data assimilation system, Q. J. R. Meteorol. Soc., 137(656), 553–597, doi:10.1002/qj.828, 2011.

Skogen, M. and Søiland, H.: A user's guide to NORWECOM v2.0. The NORWegian ECOlogical Model system., Institute of Marine Research, Bergen., 1998.

- We keep this reference since it is the most convenient document to find informations about the original model, we also refer to Aksnes (1995) which is a peer-review publication where this model is also described.

*Editor's response: The referee's original comment is a good one, but I do not think that the authors' response is sufficient. It is unclear that the question has been fully answered, particularly in relation to the deep domain. Furthermore, a PhD thesis is not a good source for this sort of information, in part because it is not necessarily easily accessible (I have not checked this) and in part because it is grey literature (i.e. has not received the same degree of scrutiny as a published paper). Far better if the authors just include some evidence in the manuscript to support their contention that 3 years is a sufficient period of time for spin up. For instance, something to clarify what "no significant drift" means and would indicate how much the state of the model is changing with time (e.g. nutrient profiles, etc.). Among other things, this would provide a published source that the authors could use in the future.*

Reply: The thesis is not very specific about the detail of the adjustments over the first

three years and mostly describe the drift of nutrients, therefore we have computed time series of basin-average nutrients from the reference run and looked at the adjustment of nutrients from it is started from climatology in 1993 to the end of the run in 2001 and included these results in the paper. We also compare to the result of the PhD thesis (which is available online:

<http://www.nersc.no/sites/www.nersc.no/files/tese%20copy.pdf>.) and continue to refer to it, the text now reads:

“In order to spin up the model, it was then run with the original parameters from 1993-1995. During the spin-up there was an adjustment of basin-averaged silicate of about  $0.2 \text{ mmol/m}^3$  during the first year, followed by a decreasing trend of about  $0.2 \text{ mmol/m}^3$  per decade that continues throughout the model run. For the basin-averaged nitrate and phosphate there are no initial adjustments, but throughout the run there are decreasing trends of less than  $0.1 \text{ mmol/m}^3$  and  $0.004 \text{ mmol/m}^3$  per decade respectively. The drift of nutrients was investigated previously in a 50-year run with HYCOM-NORWECOM V1.0 on the coarse model grid also used in this study. There was a small drift for the concentration of all three nutrients of  $\sim 0.1 \text{ mmol/m}^3$  for nitrate (increase) and silicate (decrease) and a  $\sim 0.01 \text{ mmol/m}^3$  decrease of phosphate (Hansen, 2008), therefore we do not anticipate that this drift will subside with time and conclude that three years spin-up is sufficient for the system.”

*Editor's response: Per the previous point, could it simply be that the deeper part of the domain is less perturbed by the biogeochemical model? And on the point concerning a lack of response despite parameter changes could, again, this have something to do with the short duration of the spin-up? Biogeochemical fluxes are typically smaller at depth, so longer spin-ups are can be required to determine whether the model is in "steady state". Please ensure that the text includes such caveats where appropriate.*

Reply: We have included on sentence about this in the text under section 3.3:

“Processes in the deeper layers are slower and therefor we do not expect impacts by the parameter alterations in the biogeochemical model on the time scale of the model simulation period.”

*Editor's response: Figure 9 is impenetrable. It might be preferable to show the model-observation differences as column plots in which the x-axis is month (4 columns) and the y-axis is depth (n rows), with a blue-red colour scale per Figure 5 (i.e. underestimates blue, overestimates red). This would have its own problems (not least that you'd need separate panels for the original and new simulations), but as it is very difficult to make any sense from the plot as it currently appears, an alternative is worth trying.*

Reply: We could not find a good solution where we combined all three nutrients, therefore Figure 9 now only shows nitrate, while silicate and phosphate are shown in supplementary materials. The text has been altered accordingly.

*Editor's response: Regarding Figure R1, ensure that you comment on whether such a sporadic pattern of skill / no skill could just be random, or whether it reflects real skill in the regions identified as such. Where neighbouring grid cells exhibit very different levels of skill, rather than showing a continuous gradient of skill, one suspects that skill is, in reality, low. Models can be right for the wrong reasons some of the time. Again, material added concerning spin-up*

Reply: By mistake, figure R1 shows the error statistics at depth 500-1500 meters, the

figure, the results in the 100-500 meter range is largely ‘no-skill’. As we see it there are two issues here:

- (1) Is the model efficiency pattern in R1 random? Yes, to some degree, we tested this by moving the boundaries of the boxes and in some areas the skill changes, the skill will generally be lower if larger boxes are used. For the bias, the large-scale pattern in R2 persist with changing boundaries.
- (2) Is the low skill in the model caused by drift in the model? There is a low overall drift in the model, however there is also a diffusion vertical of the nutrient gradient in the 100-1000 meter layer which will cause a lower skill in this region. However, since the model efficiency is the model error scaled by the variability in the data the most important reason for the low skill at depth is that the variability in nutrients at depth is so low that even small errors will give very poor model efficiency. This is also mention in the paper in section 3.1: “However, since the observed nutrients have low variability below 500 meters the ME shows no skill in most regions.”

*Editor’s response: First of all, were the relative sizes in the parameter changes consistent? It looks to me like grazing was altered to a much greater extent than was mortality (in percent terms). Secondly, the mortality term is described as quadratic but appears to be something different in Table 2. Thirdly, this term has an additional coefficient,  $k_6$ , whose value likely has a bearing on the insensitivity, but this is not discussed at all. Fourthly, the authors present no information at all on the behaviour of the modelled zooplankton – were they insensitive to the change, or was it just the error statistics?*

Reply: We find it difficult to compare the changes in the maximum grazing rate to the mortality because the resulting growth will also be depending on the food availability and temperature. But, yes, in terms of percentage the change was much larger for the microzooplankton grazing rate, and we also see that for microzooplankton the change in the grazing parameter caused a larger change than the mortality modifications.

Concerning the mortality in run N04, the mortality rate is linear while the resulting mortality (the amount of zooplankton lost) will be quadratic and we therefore refer to this mortality as quadratic mortality. We have checked the use of ‘mortality’ and ‘mortality rate’ and changed two instances where the wrong one was used.

In figure R4 we show both the mortality rate and the mortality in the four runs N00, N04, N05 and N06 over the range of typical zooplankton concentration. The parameter  $k_6$  would regulate how steeply the initial slope of the mortality rate increases at small zooplankton concentration, but will have a smaller effect at larger zooplankton concentrations.

There is an also an effect on the zooplankton concentration in these runs.

We have modified the discussion about sensitivity to zooplankton parameters in the text: “Several of the parameter alterations investigated had little impact on the results of the model as seen in the error statistic. Quadratic, rather than linear, mortality in the phytoplankton was one of the changes that had little effect on the error statistics, while a change in the grazing rates had a large effect. The sensitivity of this model to

the diet compositions of zooplankton has also been shown in a more theoretical study on parameter estimation by data assimilation by Simon et al. (2012). It is a factor that the change to the grazing parameter was larger than the changes to the mortality parameters, However, the changes in N04 cause the mesozooplankton to increase by about 60% and (the other alteration of the mortality rate caused changes of the same order of magnitude) and this indicates that rather large changes in the zooplankton concentrations are needed to perturb the nutrients and phytoplankton concentrations. The zooplankton mortality is the closure term in the model, but contrary to other studies (e.g Steele and Henderson, 1992) perturbations of the functional form of the mortality in N04 had little effect on the results. The reason for the lack of sensitivity of the nutrients and phytoplankton to changes in the mortality term is not clear.”

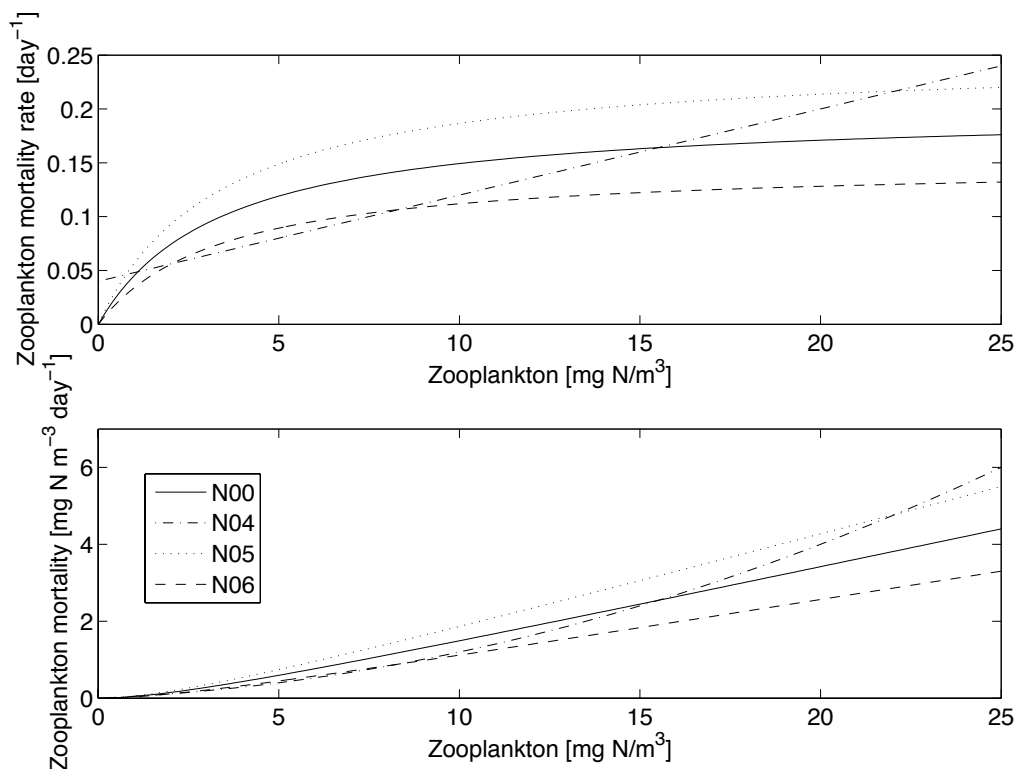


Figure R4. The mortality rate (upper) and mortality (lower) for different runs as a function of zooplankton concentration.

*Editor’s response: Perhaps the authors could consider adding such material here? It seems highly pertinent given that the authors are trying to validate the model. Note that there is scope for adding it as supplementary material should it not sit easily in the text. Adding “not shown” is not a useful response to this question.*

We have added some supplementary figures showing this.

*Editor’s response: See previous comment. Saying “not shown” is not unambiguous nor is it helpful here. Please consider adding the material you refer to as either supplementary material or directly to the manuscript. Incidentally, presumably when you say “figure 3” in your referee response you mean Figure R3? Will this be included in the final manuscript?*

Reply: I have removed the ‘not shown’ after this sentence and changed the sentence to “None of the parameter alterations affecting the timing of the spring bloom by more than a few days, while the lag in bloom initiation compared to observations is

20-30 days.” Here we refer to figure 3 in the Samuelsen, 2009 paper which clearly shows the lag in the spring bloom compared to observations, we have added this in the paper by changing the reference to (Figure 3: Samuelsen et al, 2009b).

*Editor’s response: Wearing my pedant’s hat I would rephrase this to “Biogeochemical processes in the ocean are less well understood than those of physics, and model formulations and parameterisations are correspondingly less well developed or constrained. Additionally, relative to ocean physics, there are fewer observational data for the validation and evaluation of modelled biogeochemistry.”*  
Reply: The sentences have been changed according to this suggestion

*Editor’s response: “Hereby” makes no sense; I think you mean “Here”*  
Reply: Changed

*Editor’s response: Need to pluralise to – “Analyses have shown ...”*  
Reply: Changed

*Editor’s response: Steele & Henderson (1992) were concerned with how model behaviour changed in response to different functional forms of the closure term, not strictly changes in the associated model parameters (e.g. the maximum loss rate). You seem to be conflating functional forms and parameters here. Please tighten up your wording so that it is clear what you’re referring to and whether your sensitivity test is congruent with Steele & Henderson (1992).*  
The sentence is now: “The zooplankton mortality is the closure term in the model, but contrary to other studies (e.g Steele and Henderson, 1992) perturbations of the functional form of the mortality in N04 had little effect on the results.

*Editor comment: The modification made in the text does not convey this information. I misinterpreted this statement to refer to bioconvection. Please rectify this for clarity.*  
Reply: This information was added before the reference to Backhaus in the paper.

Authors’ response: (7) We have gone through the manuscript and checked out uses of data, it has been changed in the figure labels as well.

*Editor comment: A commonly used convention is that observational science produces “data” while modelling science produces “output” (or “model data”, though this is clunky).*  
Reply: Throughout the manuscript we have used data when referring to in-situ or satellite observation. About the model output we have used ‘model simulation’ or ‘model results’. Two places ‘model data’ have been replaced by ‘model results’. We think now there is no room to misunderstand to which one we refer, which I think was the reviewers main issue with out use of these words.

*Editor comment: The code availability section usually appears towards the end of a manuscript. Please examine recently published GMD papers for guidance (I should have amended this previously).*  
Reply: we move the part about the code availability to the end of the paper.

1 **Tuning and assessment of the HYCOM-NORWECOM V2.1**  
2 **biogeochemical modeling system for the north Atlantic and**  
3 **Arctic Ocean**

4

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11

12 **Abstract**

13 The HYCOM-NORWECOM modeling system is used both for basic research and as a part of  
14 the forecasting system for the Arctic Marine Forecasting Centre through the MyOcean  
15 project. Here we present a revised version of this model. The present model, as well as the  
16 sensitivity simulations leading up to this version, have been compared to a dataset of in-situ  
17 measurements of nutrient and chlorophyll from the Norwegian Sea and the Atlantic sector of  
18 the Arctic Ocean. The model revisions having most impact included adding diatoms to the  
19 diet of micro-zooplankton, increasing micro-zooplankton grazing rate and decreased silicate-  
20 to-nitrate ratio in diatoms. Model runs are performed both with a coarse- (~50 km) and  
21 higher-resolution (~15km) model configuration, both covering the North Atlantic and Arctic  
22 Ocean. While the new model formulation improves the results in both the coarse- and high-  
23 resolution model, the nutrient bias is smaller in the high-resolution model, probably as a result  
24 of the better resolution of the main processes and improved circulation. The final revised  
25 version delivers satisfactory results for all three nutrients as well as improved result for  
26 chlorophyll in terms of the annual cycle amplitude. However, for chlorophyll the correlation  
27 with in-situ data remains relatively low. Besides the large uncertainties associated with  
28 observational data this is possibly caused by the fact that constant C/N- and CHL/N ratios are  
29 implemented in the model.

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## 1 Introduction

Physical ocean forecasting systems are now operational in many ocean regions (Le Traon, 2013) and in several forecasting systems biogeochemical models have been included (Edwards et al., 2012; Wan et al., 2012). Biogeochemical processes in the ocean are less well understood than those of physics, and model formulations and parameterizations are correspondingly less well developed or constrained. Additionally, relative to ocean physics, there are fewer observational data for the validation and evaluation of modelled biogeochemistry. At the same time, operational systems including biogeochemical variables can supply valuable information on environmental indicators such as oxygen concentration, N/P-ratios, and algae concentrations. Over time, they may give information on accumulated quantities, such as annual primary production and inter-annual variability in phytoplankton production. Data assimilation is also being used for improving the model predictions (Sakov et al., 2012) and for estimating unknown parameters, the assimilation of ocean color data in operational models is underway.

HYCOM-NORWECOM is used as a part of the operational system for the Arctic (the Arctic Marine Forecasting Centre) implemented through the EU-FP7 supported MyOcean project. The biogeochemical forecast has been operational since the fall of 2011. In connection to the setup of the biogeochemical part of the forecasting system, a series of sensitivity runs testing alternative model formulations were performed and a subsequent update of the HYCOM-NORWECOM system was implemented. The final model formulation chosen was uploaded to the forecasting system in October 2012 and is now the operational model used. Daily values of nutrient, phytoplankton, oxygen etc. can be browsed at <http://www.myocean.eu/web/24-catalogue.php> and downloaded after registration. Focal areas for this study are the Nordic Seas and the Arctic. These areas contribute to a large fraction of the world ocean carbon sink (Takahashi et al., 2009). Aside from assessing the whole model area (Fig. 1) we focus the comparison on two smaller regions, one in the Norwegian Sea, important area for the heat transport into the Nordic Seas and one in the Barents Sea where one of the branches of Atlantic Water enters the Arctic Ocean.

Here we present HYCOM-NORWECOM V2.0 and V2.1 together with the sensitivity simulations leading up to the V2.1 (Table 1). The model results are evaluated against an in-situ dataset for the Norwegian Sea and the statistical results are presented. The HYCOM-

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**Deleted:** Not all biogeochemical processes in the ocean are well understood and therefore biogeochemical models are less accurate than circulation models both with respect to model formulations and parameterizations. Observational data for validation and model evaluation are scarcer than for circulation models.



1 NORWECOM model was tested against local in-situ observations and derived gridded  
2 climatology of nutrients, as well as satellite data. However, we found that the in-situ data was  
3 the most instructive and the tuning relied most heavily on this dataset when making the  
4 upgrade. Statistical measures of the models performance for each of the parameter sets were  
5 calculated in sub-regions as well for the entire area.

## 6 **2 Methods**

### 7 **2.1 Model description**

8 HYCOM-NORWECOM is a coupled physical biological modeling system. HYCOM  
9 (v2.2.12), the HYbrid Coordinate Ocean Model (Bleck, 2002), is an ocean model using hybrid  
10 coordinates; isopycnal coordinates in the deep stratified waters, and z-level coordinates in the  
11 upper mixed layer. A description of this setup of HYCOM can be found in Sakov et al.  
12 (2012) and user guides for the different versions of HYCOM are available online at  
13 <http://hycom.org/hycom/documentation>. HYCOM is routinely used for forecasting and the  
14 predictions are regularly evaluated using in-situ and remote-sensing observations of salinity,  
15 temperature and sea ice (<http://myocean.met.no/ARC-MFC/V2Validation/index.html>).  
16 Comparisons between observations, free-runs (used in this study) and assimilative runs can be  
17 found in Sakov et al. (2012) and Samuelsen et al. (Samuelsen et al., 2009a). NORWECOM  
18 (Aksnes et al., 1995; Skogen and Søiland, 1998) is currently run with 11 variables: nitrate,  
19 phosphate, silicate, diatoms, flagellates, micro- and meso-zooplankton, nitrogen detritus,  
20 phosphorous detritus, biogenic silica and oxygen (Fig. 2). The micro- and meso-zooplankton  
21 were recently added and use the formulations and parameters defined in ECOHAM (Pätsch  
22 and Kühn, 2008; Stegert et al., 2009). The coupling of NORWECOM towards HYCOM was  
23 first done in 2005 and has been used for several studies in the Norwegian Sea and North  
24 Atlantic (Hansen et al., 2010; Samuelsen et al., 2009b). An overview of the different version  
25 can be found in Table 1.

26 The complete description of the NORWECOM V2.0 can be found in the user guide (Skogen  
27 and Søiland, 1998), below we provide a description of the differences in the biogeochemical  
28 formulations in HYCOM-NORWECOM here compared to that version. With regards to  
29 nutrient limitation the NORWECOM V2.0 applied a multiplicative relationship for the total  
30 growth ( $\mu_{phy}$ ) of phytoplankton:

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$$\mu_{phy} = \mu_{max} \times Rad\_lim \times \prod_{i=1}^n Nut\_lim_i \quad (1)$$

Where  $\mu_{max}$  is the maximum growth rate,  $Rad\_lim$  is the growth limitation due to light and  $Nut\_lim_i$  is the growth limitation for nutrient  $i$ . In HYCOM-NORWECOM it is the minimum of the limitation factors that determines the growth:

$$\mu_{phy} = \mu_{max} \times \min(Rad\_lim, Nut\_lim_{i,j=1,n}) \quad (2)$$

Except for when growth is not limited, formulation (1) will give a smaller growth rate than formulation (2) since the value of the limitation of light and nutrients are always between 0 and 1.

As in NORWECOM V2.0 (Skogen and Søliland, 1998), the main distinction between diatoms and flagellates in NORWECOM is that diatoms consume and is limited by silicate in addition to phosphate and nitrate. Diatoms have higher maximum growth rate than flagellates (Table 2), but the temperature-dependence for growth is the same, following Eppley (1972). The half saturation constants for nitrate and phosphate are smaller for flagellates ( $K_N=1.5$  mmol/m<sup>3</sup> and  $K_P=0.094$  mmol/m<sup>3</sup>) than for diatoms ( $K_N=2.0$  mmol/m<sup>3</sup> and  $K_P=0.125$  mmol/m<sup>3</sup>). The model assumes constant N/Chl-ratio (11 g N/g Chl in the control run).

NORWECOM V2.0 was primarily applied to the North Sea, while HYCOM-NORWECOM, focused the open ocean regions of the North Atlantic, therefore the extinction coefficient due to water and non-chlorophyll substances was reduced from 0.07 to 0.04 (Hansen and Samuelsen, 2009).

NORWECOM V2.0 (Skogen and Søliland, 1998) did not include zooplankton, but now there is an option of running the model with two zooplankton components, microzooplankton and mezozooplankton. The formulations for zooplankton are the same as in ECOHAM (Pätsch and Kühn, 2008), but modified to adjust for differences in the food-web structure. In HYCOM-NORWECOM, the mortality rate for phytoplankton independent of grazing is 0.035. When zooplankton is excluded, a quadratic relationship representing both grazing and other causes of mortality is used. Zooplankton grazing ( $G$ ) by a size-class of zooplankton ( $Z$ ) on a specific food source ( $fs$ ) is described by:

$$G_{fs,Z} = \frac{T_{fac}g}{k + \sum P_{fs,Z} \cdot fs} \cdot fs \cdot Z \quad (3)$$

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1 Here,  $T_{fac}$  is the temperature dependence  $T_{fac} = 1.5^{\frac{T-T_0}{10}}$ , where T is the local temperature and  
2  $T_0$  is set to 10°C, g is the maximum grazing rate (0.4 day<sup>-1</sup> for mesozooplankton and 0.5 day<sup>-1</sup>  
3 for microzooplankton) and k is the half saturation constant for zooplankton grazing which is  
4 set to 1 mmolN/m<sup>3</sup> for both size classes of zooplankton.

$$5 \quad P_{fs,z} = \frac{p_{i_{fs,z}} f s_i}{\sum p_{i_{fs,z}} f s_i} \quad (4)$$

6 where  $p_{i_{fs}}$  are the grazing preferences for the different food sources, the grazing preferences  
7 for microzooplankton can be found in Table 2, while the preferences for mesozooplankton are  
8 0.45 for diatoms and 0.275 for both microzooplankton and detritus.

9 The assimilation efficiency for both size-classes of zooplankton is set to 0.75 (Pätsch et al.,  
10 2009) and the mortality rate ( $M_z$ ) is also formulated as a half saturation relationship:

$$11 \quad M_z = m_z \frac{Z}{k_m + Z} \quad (5)$$

12 where  $m_z$  is the maximum mortality rate (0.2 day<sup>-1</sup>) and the half saturation constant  $k_m$  is 0.2  
13 mmolN/m<sup>3</sup> for both size classes of zooplankton. For the loss terms of zooplankton 90% of  
14 the material goes into the detritus pool and 10% is returned to nitrate.

## 15 2.2 Experiment setup

16 The tuning was done on a coarser grid (30-50 km) than the 15-km grid (Fig. 1) used in the  
17 operational runs to limit the computational cost, as the 15-km model takes about 5 times as  
18 long to run. The model was forced by the ERA-Interim (Dee et al., 2011) from 1989 and  
19 ERA40 (Uppala et al., 2005) for the period prior to 1989 (only spinup). The physical model  
20 was initialized from rest with climatological temperatures and salinity from the GDEM  
21 (Carnes, 2009). The biogeochemical model was initialized from climatological nutrients and  
22 oxygen values from the World's Ocean Atlas (WOA2001: Conkright et al., 2002) and constant  
23 low values for the other variables in 1993. Throughout the run relaxation back to  
24 climatological temperature, salinity, nutrients and oxygen was applied at the lateral  
25 boundaries. A weak relaxation of salinity (relaxation timescale of 200 days) was also applied  
26 at the surface. River nutrients were derived from GlobalNEWS model output (Seitzinger et  
27 al., 2005). In all, 16 sensitivity simulations were performed with the coarse model  
28 (simulation names starting with N) and the parameter changes in each run are summarized in

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1 Table 2 and the location of the relevant code is given in Table A1. In order to assess the  
2 effect of the revised parameter set on the 15-km model, two simulations were performed; one  
3 with the with the higher resolved grid (simulation names starting with TP); the original set of  
4 parameters (TP0) and one with revised set of parameters (TP1). The model was started from  
5 climatological nutrient values and constant low values for the other variables in 1993. In  
6 order to spin up the model, it was then run with the original parameters from 1993-1995.  
7 During the spin-up there was an adjustment of basin-averaged silicate of about 0.2 mmol/m<sup>3</sup>  
8 during the first year, followed by a decreasing trend of about 0.2 mmol/m<sup>3</sup> per decade that  
9 continues throughout the model run. For the basin-averaged nitrate and phosphate there are  
10 no initial adjustments, but throughout the run there are decreasing trends of less than 0.1  
11 mmol/m<sup>3</sup> and 0.004 mmol/m<sup>3</sup> per decade respectively. The drift of nutrients was investigated  
12 previously in a 50-year run with HYCOM-NORWECOM V1.0 on the coarse model grid also  
13 used in this study. There was a small drift for the concentration of all three nutrients of ~0.1  
14 mmol/m<sup>3</sup> for nitrate (increase) and silicate (decrease) and a ~0.01 mmol/m<sup>3</sup> decrease of  
15 phosphate (Hansen, 2008), therefore we do not anticipate that this drift will subside with time  
16 and conclude that three years spin-up is sufficient for the system. The sensitivity simulations  
17 were initiated in 1996 and run for a 6-year period. The impact of a single parameter or model  
18 formulation change was investigated in 11 sensitivity simulations. Subsequently the impact  
19 of five different combinations of these alterations was studied. Model-observation  
20 comparisons were performed in the period 1998 to 2001 because of relatively good in-situ  
21 data coverage combined with availability of ocean colour data in this period.

22 The model results to be compared to in-situ data was extracted from the model from files  
23 containing daily averages. The modeled values from the grid box and model layer containing  
24 the observation point on the day of the observation were selected. The model results were not  
25 interpolated temporally or spatially. In the case of several observations within the same grid  
26 cell and layer, the mean of the observed values was used.

### 27 2.3 Description of observations

28 An observational dataset collected as a part of the Norwegian Institute of Marine Research  
29 monitoring activities was used. In addition to comparing the simulations to the entire dataset,  
30 we also focused the comparison on two sub-regions; one in the Norwegian Sea and one the  
31 Barents Sea (Fig. 3). The available in-situ data relevant to the NORWECOM model are  
32 nutrients (silicate, nitrate, nitrite and phosphate) and chlorophyll, obtained by analysis of

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1 discrete water samples. Because we only have one type of nitrogen nutrient source in the  
2 model, the modeled nitrate was compared to the sum of observed nitrate and nitrite. The  
3 Norwegian Sea sub-region includes Station M and thus observational data are available  
4 throughout the year for all of the variables, while in the Barents Sea observations are collected  
5 primarily during August and September (Fig. 3).

## 6 **2.4 Statistical method for model evaluation**

7 In the paper by Allen et al. (2007), several metrics for evaluation of biogeochemical models  
8 were presented. A combination of model efficiency (ME) and percentage model bias (Pbias)  
9 was used for the comparison between the model simulations and observations. These  
10 statistical quantities are defined as:

$$11 \quad ME = 1 - \frac{\sum_{n=1}^N (D_n - M_n)^2}{\sum_{n=1}^N (D_n - \bar{D})^2} \quad (6)$$

12 where  $D_n$  is observation from station  $n$ ,  $M_n$  is the corresponding model estimate,  $\bar{D}$  is the  
13 mean of the observations, and  $N$  is the total number of stations. The model efficiency is a  
14 measure of the model-observation misfit in relation to the variability of the observational data.

$$15 \quad Pbias = \frac{\sum_{n=1}^N (D_n - M_n)}{\sum_{n=1}^N D_n} \times 100 \quad (7)$$

16 Pbias gives an indication on whether the model results are consistently under- or  
17 overestimated compared to the observations.

18 In addition, standard deviation, correlation coefficient and the centered root mean square error  
19 of chlorophyll and nutrients were evaluated in Taylor diagrams (Taylor, 2001) that show the  
20 overall quality of the runs.

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## 1 3 Results

### 2 3.1 Performance of control runs

3 The model efficiency showed that the results from the control runs with the original  
4 parameters (N00 and TP0) were in general good with respect to nutrients (Fig. 4). The model  
5 performance was better for nitrate and phosphate than for silicate. In terms of ME for the  
6 nutrients there is little difference between the coarse and the fine model, but the results from  
7 the high-resolution model is slightly better. The percentage bias is also similar in the two  
8 control runs and again the estimates of nitrate and phosphate have higher skill compared to  
9 silicate (Fig. 5). The bias is positive, meaning that the modeled nutrients are consistently  
10 lower than the observed nutrients (eq. 7). The nutrient bias is slightly better in the high-  
11 resolution model than the coarse model. Below 500 meters (not shown), nitrate and  
12 phosphate are generally excellent in terms of bias, while silicate varies from excellent to  
13 good, except for a region in the central Norwegian Sea where it is poor. However, since the  
14 observed nutrients have low variability below 500 meters the ME shows no skill in most  
15 regions. Below 500 meters the model is probably quite influenced by both initial condition  
16 and the relaxation towards climatological nutrients at the boundary, as the residence time for  
17 the deep waters is estimated to be 2-10 years (Aagaard et al., 1985). Above 500 meters, the  
18 biases are generally poorer, while the model shows some skill in terms of predicting the  
19 observed nutrients. For the upper waters masses the residence time in this region it is about 3  
20 month (Poulain et al., 1996), hence the initial and boundary condition have limited influence  
21 there.

22 The prediction of the chlorophyll content is even more challenging than for the nutrients.  
23 Here, the runs with the original parameter set for both resolutions show no skill for the ME  
24 (Fig. 4) and large negative percentage biases (Fig. 5), meaning that the model consistently  
25 overestimates the chlorophyll. For chlorophyll there is no consistent improvement with  
26 resolution. Correlation between the observed and modeled chlorophyll is poor and the  
27 amplitude of the annual cycle is overestimated (Fig. 6). Analyses have shown that the model  
28 runs are consistently late in the spring bloom, a persistent feature in this model system (Figure  
29 3: Samuelsen et al., 2009b).

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### 1 3.2 Parameter modifications

2 As seen in section 3.1, the main challenge of the model lies in the overestimation of  
3 chlorophyll during the summer months. Many of the parameter changes were thus aimed at  
4 reducing the error in the phytoplankton fields, but as seen in figures 4 and 5 many of the  
5 changes had a positive influence on the simulated nutrient values as well. The original and  
6 new model formulations and parameter values of all the sensitivity simulations are listed in  
7 Table 2.

8 The first run, N01, had quadratic rather than linear mortality of phytoplankton, this change  
9 was aimed at increasing the phytoplankton losses during periods with high phytoplankton  
10 biomass. This alteration had little effect on the results, nevertheless it was also tried in  
11 combination with other parameter changes, N07 and N13, but no improvement was observed,  
12 therefore this alteration was not included in the final model formulation.

13 In nature, a wide range of Si:N ratios are observed in diatoms (Sarhou et al., 2005), therefore  
14 the second and third run, N02 and N03, altered the fixed uptake ratio of Si:N for diatoms, by  
15 decreasing and increasing this value by 25% respectively. In the control runs the model  
16 tended to consume all the silicate before nitrate in the spring, while this was not the case in  
17 the observations. A reduction in this ratio improved the modeled silicate in terms of model  
18 efficiency, while estimates of nitrate and phosphate gets reduced skill. This change however,  
19 reduced the summer chlorophyll concentrations, most likely because the spring diatom bloom  
20 consumed more nitrate, which is the limiting nutrient during the summer bloom. Increasing  
21 the ratio had the opposite effect. Because large flagellate summer concentration has been a  
22 recurring challenge in the model the reduced Si:N ratio was retained in some of the  
23 subsequent runs.

24 The next three sensitivity simulations explored alterations to the zooplankton mortality term;  
25 quadratic mortality (for both zooplankton size classes) – N04, increased and decreased  
26 mesozooplankton mortality – N05 and N06. These alterations had little effect on the error  
27 statistics and were not considered in any of the subsequent runs.

28 Three runs where the sensitivity to the choice of nitrate to chlorophyll ratio was investigated.  
29 The first (N08) was a simple increase by 25%, while the values of 12.5 (N09) and 6.3 (N10)  
30 were found in the literature (Fouilland et al., 2007; Yentsch and Vaccaro, 1958). In the North  
31 Atlantic values varying from 1 to 12.5 was found in the literature (Fouilland et al., 2007;  
32 Yentsch and Vaccaro, 1958). The alteration had little effect on the overall results for nutrient,

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1 but a rather large effect on chlorophyll. In general an increase of this ratio lead to an  
2 improvement in the chlorophyll comparison and a decrease to deterioration of the model  
3 results. We did not alter this value during the tuning, but think that a mechanistic model  
4 allowing for variable N:Chl ratio should be included in the model.

5 Motivated by the observation that diatoms can be consumed by microzooplankton (Sarhou et  
6 al., 2005) we made an experiment where diatoms were included in the diet of  
7 microzooplankton (N11). The microzooplankton grazing rate was also increased (N12).  
8 These runs, especially N12, had a negative effect on the silicate results, but a positive effect  
9 on the nitrate and phosphate. These changes also contributed to better results for the  
10 chlorophyll. The increased microzooplankton grazing rate resulted in improved performance  
11 of the model and it was the first simulation where the biases in both 1998 and 1999 were  
12 better than 'Poor' for chlorophyll.

13 From the above simulations we learned that reduction of the Si:N-ratio and microzooplankton  
14 grazing were the changes having the most positive impact on the model performance. Since  
15 these changes to zooplankton grazing negatively affected the silicate results, this alteration  
16 was combined with the reduction of the Si:N ratio in simulations N14 and N15. The run  
17 including diatoms in the microzooplankton diet was combined with reduced Si:N ratio in run  
18 N14, this only improved the silicate results. When these changes were also combined with  
19 increased microzooplankton grazing (N15) the results for all nutrients improved. In the last  
20 experiment, N16, a reduction of the maximum growth rate for both types of phytoplankton  
21 were added to N15, this had an additional positive effect on the chlorophyll errors. The  
22 parameter set in N16 was decided upon and studied in the high-resolution model.

### 23 **3.3 Assessment of revised model simulation**

24 The observations in some regions such as Station M and in the repeated sections (visible in  
25 the winter panel of Fig. 3) are collected more systematically and are more numerous than in  
26 the other regions. In the Norwegian Sea at Station M observations are available throughout  
27 the year, in the repeated sections each season is sampled, and an extensive survey in of the  
28 Barents Sea is done annually in August/September (Fig. 3). This should be kept in mind  
29 when comparing the performance of the run with original and revised parameters in different  
30 regions (Figs. 7 and 8). Overall the regional estimates were worse than the one including all  
31 observational data, but there are also areas where there are significant improvements. The



1 results show that in terms of Pbias, nitrate and phosphate were improved in the central  
2 Norwegian Sea and Eastern part of the Barents Sea (Fig. 7). In the northwest of the  
3 Norwegian Sea eastern part of the Barents Sea there is little improvement, but the two latter  
4 regions only have data in specific seasons (Fig. 3). For silicate the regions where there is  
5 improvement is more intermittent, but the bias in the original run was 'poor' over most of the  
6 region, this is no longer the case. The bias for chlorophyll changes sign, but not show any  
7 regional improvement. The model efficiency shows improvement in the estimates of all three  
8 nutrients, in particular in the central Norwegian Sea where the results were initially not so  
9 good (Fig. 8). Chlorophyll remains below 'no skill' in the most of the domain, except for a  
10 few places in east and north part of the domain, where it is 'good' (Fig. 8). Most of the  
11 differences between the two runs occur in the upper 100 meters. Processes in the deeper  
12 layers are slower and therefor we do not expect impacts by the parameter alterations in the  
13 biogeochemical model on the time scale of the model simulation period. The difference  
14 between the original and revised model run in the Norwegian and Barents Sea (boxes in Fig.  
15 3) in terms of chlorophyll is summarized in a Taylor diagram (Fig. 6). This Taylor diagram  
16 shows that overall the new runs are in better agreement with the observations, the  
17 improvement is mostly in terms of reduced standard error (green dashed curves). The  
18 amplitude is improved in the Norwegian Sea, but for the comparison to all observations it is  
19 now too low. There are only small differences in the correlation coefficients, but they are  
20 overall slightly lower in the run with revised parameterizations.

21 To assess the revised run at different depths, profiles in the upper 1000 meters of the water  
22 column in the Norwegian Sea box have been compared to in-situ data for nitrate, and  
23 chlorophyll (Figs. 9 and 10). Below 200 meters the differences from observations are similar  
24 for the two parameter sets. The same is the case for the upper 200 meters, during January and  
25 April when the water column is well mixed and the surface concentrations reflect the deep  
26 concentrations. During July the run with revised parameters is closer to the observation for  
27 nitrate, but further from the observations for silicate (Figure S3), during October both of these  
28 nutrients are closer to the observation with the revised parameters. For phosphate (Figure S4)  
29 the original run is close to the observations at the surface, but closer to the observations  
30 around 50 meters and also in the depth interval 400-800 meters. However, we have seen  
31 before that there is an overall improvement in the surface nutrients for the run with the revised  
32 model (Figs. 7 and 8). For chlorophyll (Fig. 10), it is clear that the overestimation of values  
33 that occurs with the original parameterization has now been reduced to give reasonable

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1 values. In April there is a clear indication in the observations that nutrients are being  
2 consumed in the upper layers, this is not the case in either of the model runs, and consistent  
3 with the modeled surface chlorophyll values that are lower than observed in this period (not  
4 shown). The late onset of the spring bloom has been a persistent challenge in the model for  
5 several years and seems to be related to delayed onset of stratification in the physical model  
6 fields, rather than the biological formulations (Samuelsen et al., 2009b).

7

## 8 **4 Discussion**

### 9 **4.1 Uncertainties connected to observations**

10 In general, the representativity of the measurements depends on how often it is measured –  
11 i.e. the uncertainty decreases with increasing number of observations. Depending on the  
12 issues addressed, there will be different requirements for geographical coverage, number of  
13 stations, frequency and parameters measured (Figure C1 - Ottersen et al., 1998).

14 Actual programs on *in situ* monitoring of the biogeochemical environment are mainly carried  
15 out by discrete sampling and subsequent analysis along with regularly monitoring cruises or  
16 by stationary measuring systems like buoys. Monitoring cruises are restricted in spatial and  
17 temporal coverage, hence limiting the availability of high quality observational data. In  
18 addition the measurement methodologies are, especially for the biogeochemical parameters,  
19 an issue in terms of uncertainty of the specific measurement (i.e. Proctor and Roesler, 2010).

20 Exemplary for the variety of biogeochemical measurements are the challenges connected to  
21 the measurements of Chl *a* concentration, which are performed by analysing filtered water  
22 samples with spectrophotometric or high-performance liquid chromatography (HPLC)  
23 methodologies which are cost intensive. In order to lower the costs, a range of autonomous  
24 sensors has been developed to overcome these limitations. These sensors measure the Chl *a*  
25 fluorescence, which is used to provide an estimate of the Chl *a* concentration.

26 In addition, when comparing to model results there is an added uncertainty in what the  
27 observations represent. One measurement may represent the value in a few litres of water,  
28 while the model value represents the value in  $\sim 10^9$  m<sup>3</sup> of water, depending on the model  
29 resolution. Here, the same dataset was used for evaluation of the effect of the tuning, as was  
30 used to study the needs for tuning. To be fully validated, the model should be compared to

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1 independent observed data (Stow et al., 2009). However, due to scarce availability of  
2 observed data, it was decided to use all data for both activities.

3

#### 4 4.2 Parameter changes

5 Most of the parameter changes were included to reduce the systematic overestimation of  
6 phytoplankton biomass during summer. Some parameter alterations were conducted to study  
7 the sensitivity of the model to the variety of ecosystem properties reported in the literature,  
8 this included different Si:N ratios and the inclusion of diatoms in the diet of  
9 microzooplankton. Several of the parameter alterations investigated had little impact on the  
10 results of the model as seen in the error statistic. Quadratic, rather than linear, mortality in the  
11 phytoplankton was one of the changes that had little effect on the error statistics, while a  
12 change in the grazing rates had a large effect. The sensitivity of this model to the diet  
13 compositions of zooplankton has also been shown in a more theoretical study on parameter  
14 estimation by data assimilation by Simon et al. (2012). It is a factor that the change to the  
15 grazing parameter was larger than the changes to the mortality parameters, However, the  
16 changes in N04 cause the mesozooplankton to increase by about 60% and (the other alteration  
17 of the mortality rate caused changes of the same order of magnitude) and this indicates that  
18 rather large changes in the zooplankton concentrations are needed to perturb the nutrients and  
19 phytoplankton concentrations. The zooplankton mortality is the closure term in the model, but  
20 contrary to other studies (e.g Steele and Henderson, 1992) perturbations of the functional  
21 form of the mortality in N04 had little effect on the results. The reason for the lack of  
22 sensitivity of the nutrients and phytoplankton to changes in the mortality term is not clear.

23 Increasing the N:Chl ratio would on one hand decrease the amount of chlorophyll per  
24 phytoplankton biomass, but also how quickly light is attenuated with depth. This alters the  
25 vertical distribution of phytoplankton, but it changes the concentrations only by a few percent,  
26 hence this effect is small compared to the effect on the chlorophyll concentration from  
27 altering the N:Chl ratio. The change of N:Chl (which is proportional to the C:Chl ratio in this  
28 model) with light availability is now well established (Geider, 1987) and implementing a  
29 variable N:Chl ratio is one of the future developments planned for this model.

30 The changes in the uptake ratio of silicate to nitrate had a large influence on the progress of  
31 both the diatom bloom and the flagellate bloom. Silicate is the limiting nutrient for diatoms,

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1 and when lowering this ratio more nitrate can be consumed leaving less nitrate for the  
2 flagellates and limiting the size of the bloom. Observed uptake ratios of Si:N vary widely and  
3 probably also varies between species, regions and seasons. Ideally a flexible uptake ratio  
4 could be included, for example as in the ERSEM model (i.e. Blackford et al., 2004), but  
5 including variable stoichiometry also increases the number of variables that has to be  
6 advected in the model and hence the computations cost considerably.

7 Because of computational limitations, only a small subset of the parameters was tested in this  
8 tuning exercise, the parameters were picked based upon past experience with the model. As  
9 grazing seems to be an important control mechanism in the model, the zooplankton  
10 assimilation efficiency may be an important parameter to test in the future. The temperature  
11 dependence of growth and respiration for both zooplankton and phytoplankton would  
12 probably influence the progress of the blooms across regions, but past experience with the  
13 model has shown that this model has little sensitivity to parameters related to phytoplankton  
14 growth, hence these parameters have been mostly left unchanged in this study. Additionally  
15 the sinking rates for detritus influence the amount of regenerated nutrients during summer.

#### 16 **4.3 Regional differences in performance**

17 Evaluating the final run (TP1) compared to all observational data (Figs. 4 and 5) and to  
18 observations in different regions (Figs, 7 and 8), it is clear that the model performed better  
19 overall than on a region-by-region basis. The explanation for this may lie partly in the  
20 placement of water masses in the model combined with the locations of the measurements. In  
21 the Norwegian Sea the majority of measurements are taken at a single location (Station M).  
22 For the model to perform well there, it needs to simulate the correct water masses at this exact  
23 point. Station M is located close to a front between two water masses, and the model is not  
24 always simulating the location of this front well (Figure S1). In the Barents Sea most of the  
25 observations are collected in sections or over the whole area during early fall, therefore some  
26 of the dependency on simulating the correct location of fronts falls away in this region. In  
27 shallow areas, such as along the coast and in the Barents Sea, better representation of benthic  
28 processes as well as the lack of tides are probably sources of errors.

29 The location of the ice edge affect the results of the biogeochemical model (Samuelsen et al.,  
30 2009a). The observations used here are primarily from open-ocean regions, so we have  
31 limited knowledge of the model performance close to the ice edge. The comparison of the

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1 physical model simulation (free-run) to satellite observations shows that the ice-edge follows  
2 the observed pattern (Sakov et al. 2012), but of course it is not 100% accurate. In the model  
3 light does not propagate through ice, and the ice edges also influences mixing, therefore errors  
4 are expected in both chlorophyll and nutrients if the model places the ice edge incorrectly.. In  
5 addition, the fact that we don't include ice-algae in the model also introduces sources of  
6 errors.

## 7 **5 Conclusions**

8 In total 18 sensitivity runs were performed on the higher- and coarser resolution model grid.  
9 First, the effect of tuning of single parameters was studied. Subsequently, the tuning of  
10 combinations of parameters were tested in the coarse model. The conclusion was that the best  
11 overall results were obtained when a combination of grazing preference for  
12 microzooplankton, Si:N ratio in diatoms and reduced growth rate for phytoplankton was used.  
13 This combination of parameters was then changed in the higher-resolution model and the  
14 differences in performance between the two sets of parameters were investigated in that  
15 configuration.

16 The revised run shows a clear improvement compared to the original run, particularly for  
17 nutrients but also for chlorophyll, but while the previous run tended to overestimate the  
18 annual cycle of chlorophyll, the revised run tends to underestimate the amplitude (Fig. 6).  
19 Based on these results, the revised parameter set presented here were also implemented as part  
20 of an operational system for the Arctic. A major difference between the model runs presented  
21 here and the operational system is that the operational system includes data assimilation in the  
22 physical model (Sakov et al., 2012), which may alter the physical model and in turn alter the  
23 performance of NORWECOM. A study of the impact of data assimilation on this model  
24 (Samuelsen et al., 2009a) showed that there were typically a difference of 5-10% for the  
25 nutrients and chlorophyll between the free run and the run with assimilation, but with  
26 difference up to 20% in the Arctic. Data assimilation can also be applied to the  
27 biogeochemical model, both as a mean of improving the forecast fields and as a method for  
28 optimizing model parameters (Simon et al., 2012).

29 We have shown that the model reproduces a reasonable annual cycle, but one persistent  
30 challenge the initiation time of the spring bloom is later than the observations. None of the  
31 parameter alterations, affecting the timing of the spring bloom by more than a few days, while  
32 the lag in bloom initiation compared to observations is 20-30 days, This indicates that the

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1 error in timing is an effect either of the physical model or a missing process, such as for  
2 example phyto-convection (the early seeding of the spring bloom by phytoplankton that was  
3 mixed down during winter; Backhaus et al., 1999). Another challenge is to show that the  
4 model also produces realistic interannual variability. The model shows less variability than  
5 the observed data, but this is also expected as the observations include a spatial and temporal  
6 variability that cannot be resolved of a model of this resolution.

7 During the tuning process the parameter sensitivity of the module was explored and the  
8 changes that were motivated by observation-based findings, for example that Si:N is highly  
9 variable and that microzooplankton are grazing on diatoms, had a positive influence on the  
10 model. This suggests that greater refinement of the models in general should be done in closer  
11 collaboration with ecologist and field oceanographers.

## 12 **Code availability**

13 The full model code is available at  
14 [https://svn.nersc.no/hycom/browser/HYCOM\\_2.2.12/CodeOnly/src\\_2.2.12/](https://svn.nersc.no/hycom/browser/HYCOM_2.2.12/CodeOnly/src_2.2.12/). The code is  
15 continually under development and version control is used when updating the code, so the  
16 HYCOM-NORWECOM V2.0 used for in the reference run, which were performed in  
17 October 2011 is revision number 186, while HYCOM-NORWECOM V2.1 corresponds  
18 revision number 224.

## 20 **Acknowledgements**

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1 Table 1. Model versions and references.

HYCOM	NORWECOM	HYCOM-NORWECOM	References
V2.2.12	V2.0	V1.0	Description:(Skogen and Søiland, 1998);  Examples of application: (Hansen and Samuelsen, 2009; Hansen et al., 2010)
V2.2.12	V2.0+zooplankton	V2.0	Application: Samuelsen and Bertino, 2011
V2.2.12	V2.0+zooplankton+ parameter tuning	V2.1	This paper

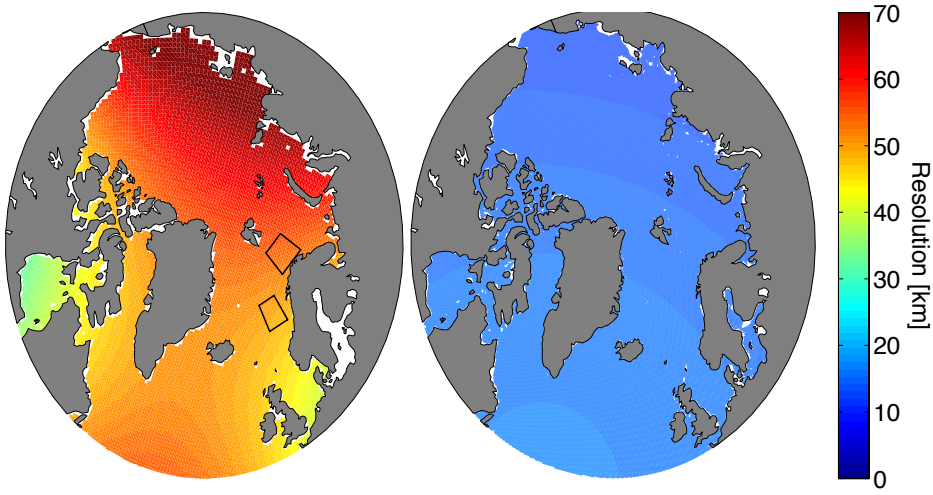
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1 Table 2. Overview of runs performed with the associated parameter values.

	Parameter for tuning	Original value	New value
N00	Reference run		
	Reference run with TPO high resolution		
N01	Quadratic mortality for phytoplankton	$cc(3), cc(3)=4.0e-7$	$cc(3)/15.0+cc(3)*P/15.0$
N02	Si:N-ratio in diatoms	$1.75 \text{ mgSi/mgN}=0.875 \text{ mmolSi/mmolN}$	$0.575 \text{ mmolSi/mmolN}=1.15 \text{ mgSi/mgN}$
N03	Si:N-ratio in diatoms	$1.75 \text{ mgSi/mgN}=0.875 \text{ mmolSi/mmolN}$	$1.175 \text{ mmolSi/mmolN}=2.35 \text{ mgSi/mgN}$
N04	Quadratic mortality in zooplankton	$m_z*(z/(z+cnit*k6)), m_z=0.2,$ $z=\text{zooplankton-conc} [\text{mgN/m}^3],$ $cnit=14.01 \text{ mgN/mmolN},$ $k6=0.2$	$m_z/5.0+m_z*z/25.0$
N05	Mesozooplankton mortality (+25%)	$m_{z\text{-meso}}=0.2$	$m_{z\text{-meso}}=0.25$
N06	Mesozooplankton mortality (-25%)	$m_{z\text{-meso}}=0.2$	$m_{z\text{-meso}}=0.15$
N07	Combination of N01 and N02	$cc(3), cc(3)=4.0e-7,$ $1.75 \text{ mgSi/mgN}$	$cc(3)/15.0+cc(3)*P/15.0,$ $1.15 \text{ mgSi/mgN}$
N08	N:Chl-ratio		11
N09	N:Chl-ratio		13.75
N10	N:Chl-ratio		12.5
N11	Grazing preferences for microzooplankton	$pi21=0.633\text{-flagellates},$ $pi24=0.367\text{-detritus}$	$pi21=0.333\text{-flagellates},$ $pi23=0.333\text{-diatoms},$ $pi24=0.333\text{-detritus}$

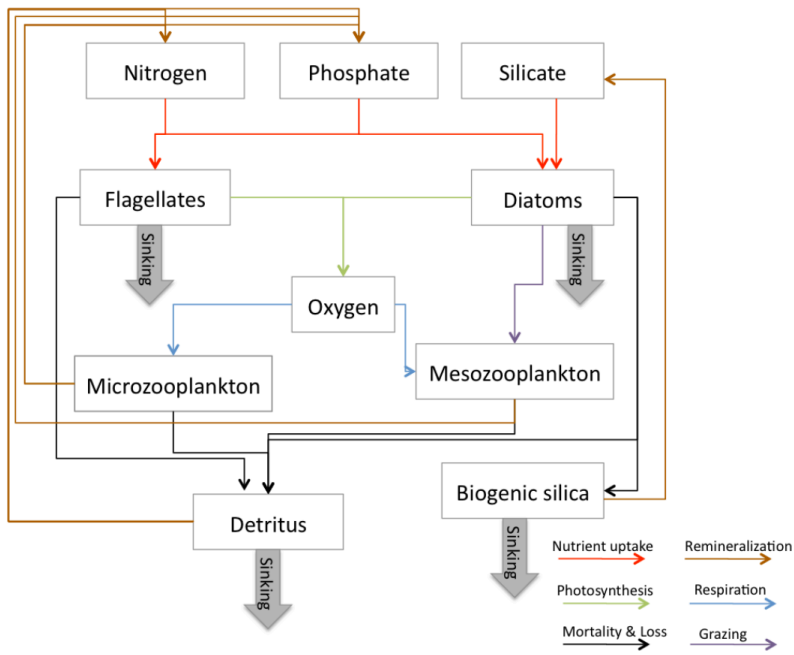
N12	Maximum microzooplankton grazing rate	g=0.5	g=1.0
N13	Combination of N11 and N1	pi21=0.633-flagellates, pi24=0.367-detritus cc(3), cc(3)=4.0e-7	pi21=0.334-flagellates, pi23=0.333-diatoms, pi24=0.333-detritus, cc(3)/15.0+cc(3)*P/15.0
N14	Combination of N11 and N2	pi21=0.633-flagellates, pi24=0.367-detritus, 1.75 mgSi/mgN	pi21=0.334-flagellates, pi23=0.333-diatomes, pi24=0.333-detritus, 1.15 mgSi/mgN
N15	Combination of N14 and N12	pi21=0.633-flagellates, pi24=0.367-detritus, 1.75 mgSi/mgN, g(micro)=0.5	pi21=0.334-flagellates, pi23=0.333-diatoms, pi24=0.333-detritus, 1.15 mgSi/mgN, g(micro)=1.0
N16	Combination of N14 and reduced growth rate for phytoplankton	pi21=0.633-flagellates, pi24=0.367-detritus, 1.75 mgSi/mgN, Vmax(dia)=1.53E-5, Vmax(fla)=1.02E-5	pi21=0.334-flagellates, pi23=0.333-diatoms, pi24=0.333-detritus, 1.15 mgSi/mgN, Vmax(dia)=1.15E-5, Vmax(fla)=0.76E-5
TP1	High-resolution run with the parameter values of N16	pi21=0.633-flagellates, pi24=0.367-detritus, 1.75 mgSi/mgN, Vmax(dia)=1.53E-5, Vmax(fla)=1.02E-5	pi21=0.334-flagellates, pi21=0.333-diatoms, pi24=0.333-detritus, 1.15 mgSi/mgN, Vmax(dia)=1.15E-5, Vmax(fla)=0.76E-5

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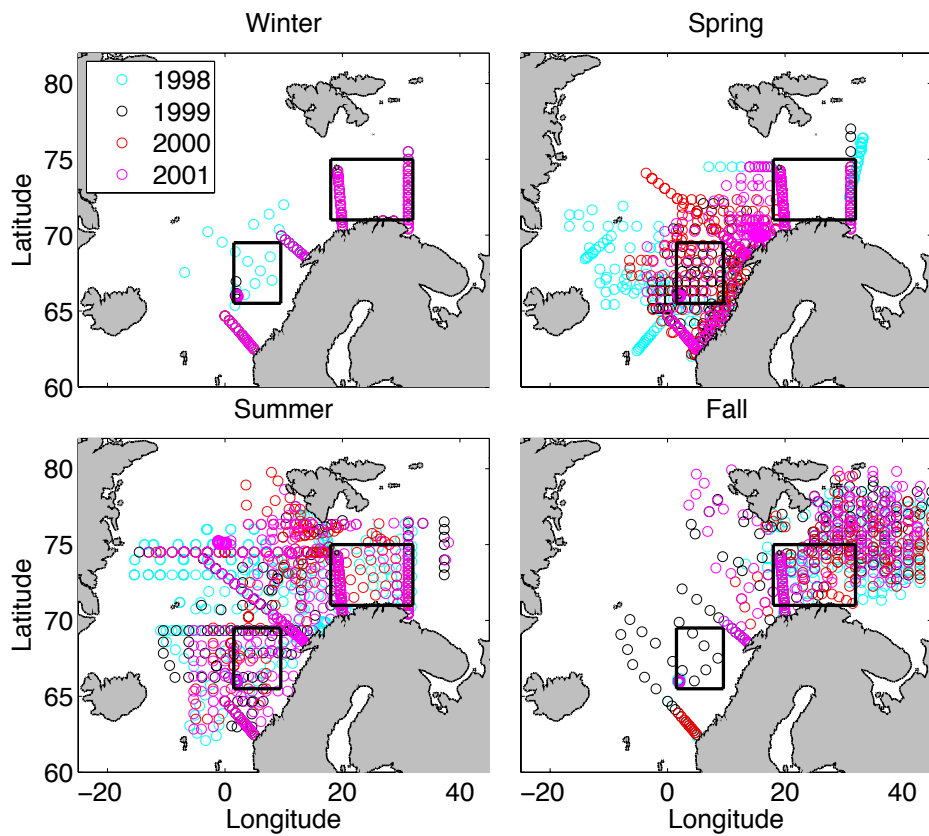
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Figure 1. Resolution of the two model grids used in this study. The two areas indicated by black lines in the map to the left are the areas referred to as Norwegian Sea – southern area - and Barents Sea – northern area.



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Figure 2. Flow chart of the interaction between the individual model components in NORWECOM.



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3 Figure 3. Spatial in-situ data coverage for nitrate in different years and seasons for the dataset  
 4 used. The coverage for the other variables is similar. The southern areas are mostly sampled  
 5 in spring and summer, while the Arctic regions are more sampled in summer and fall. There  
 6 are very few open-ocean measurements during winter, but in the sections visible in the  
 7 winter-panel (upper, left) there are observations for all years and seasons.

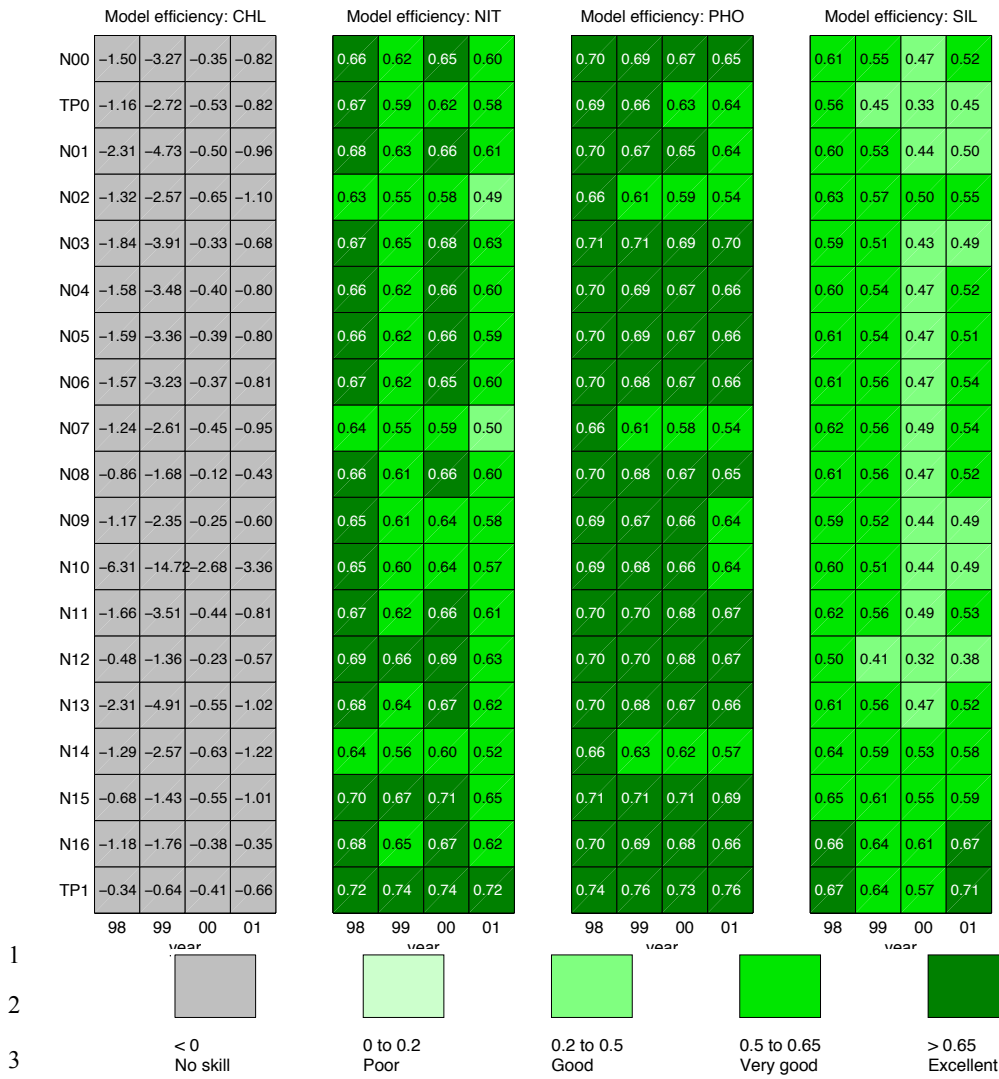


Figure 4. Model efficiency (ME, see text) for the model simulations compared to all available observations from the period 1998-2001.



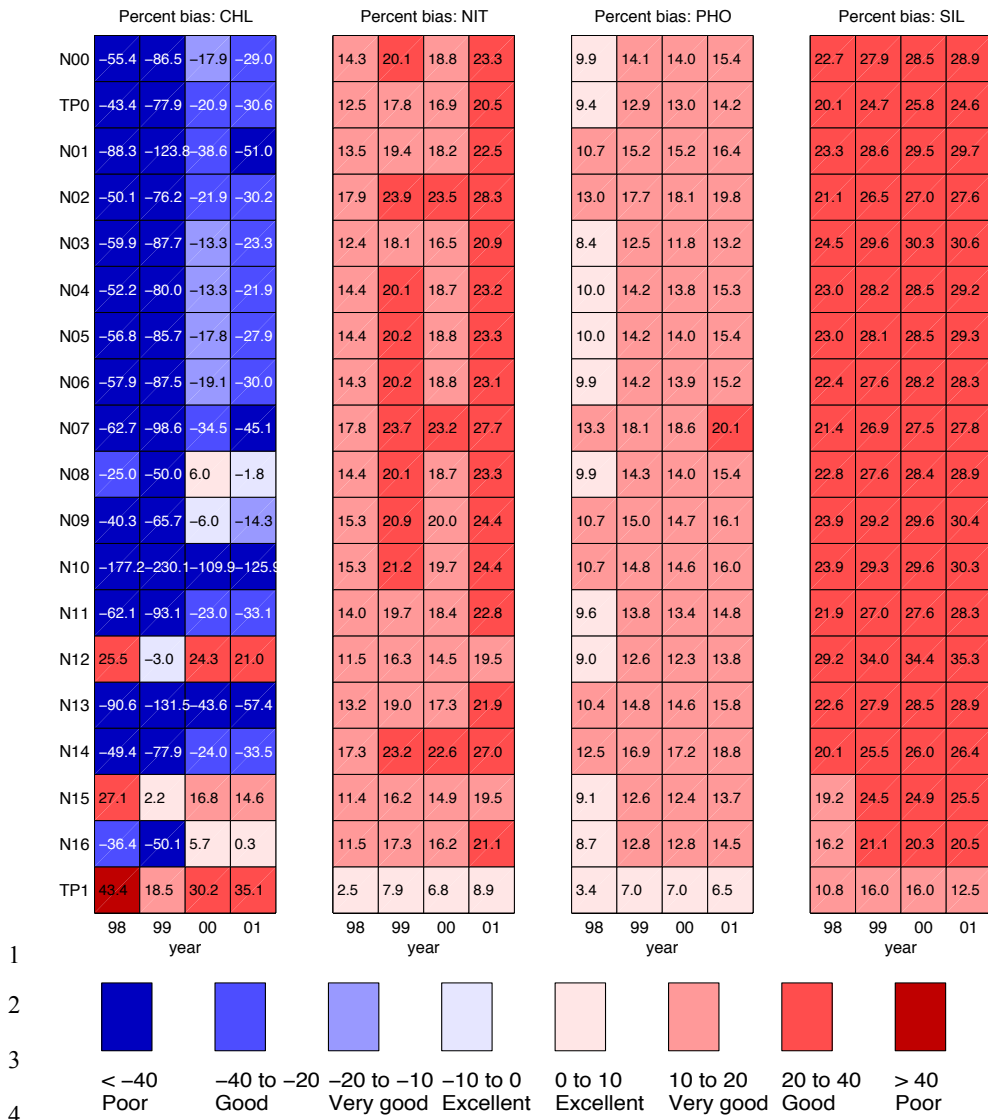
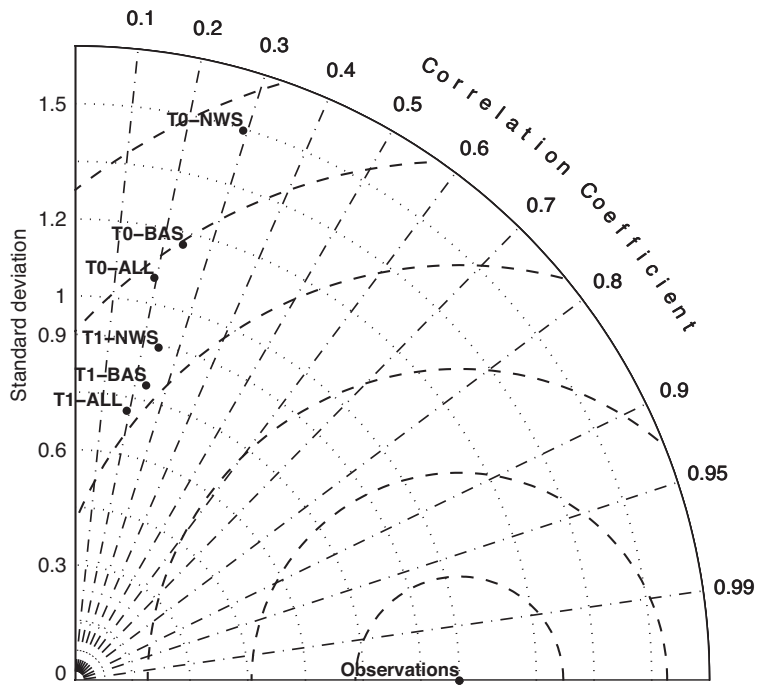


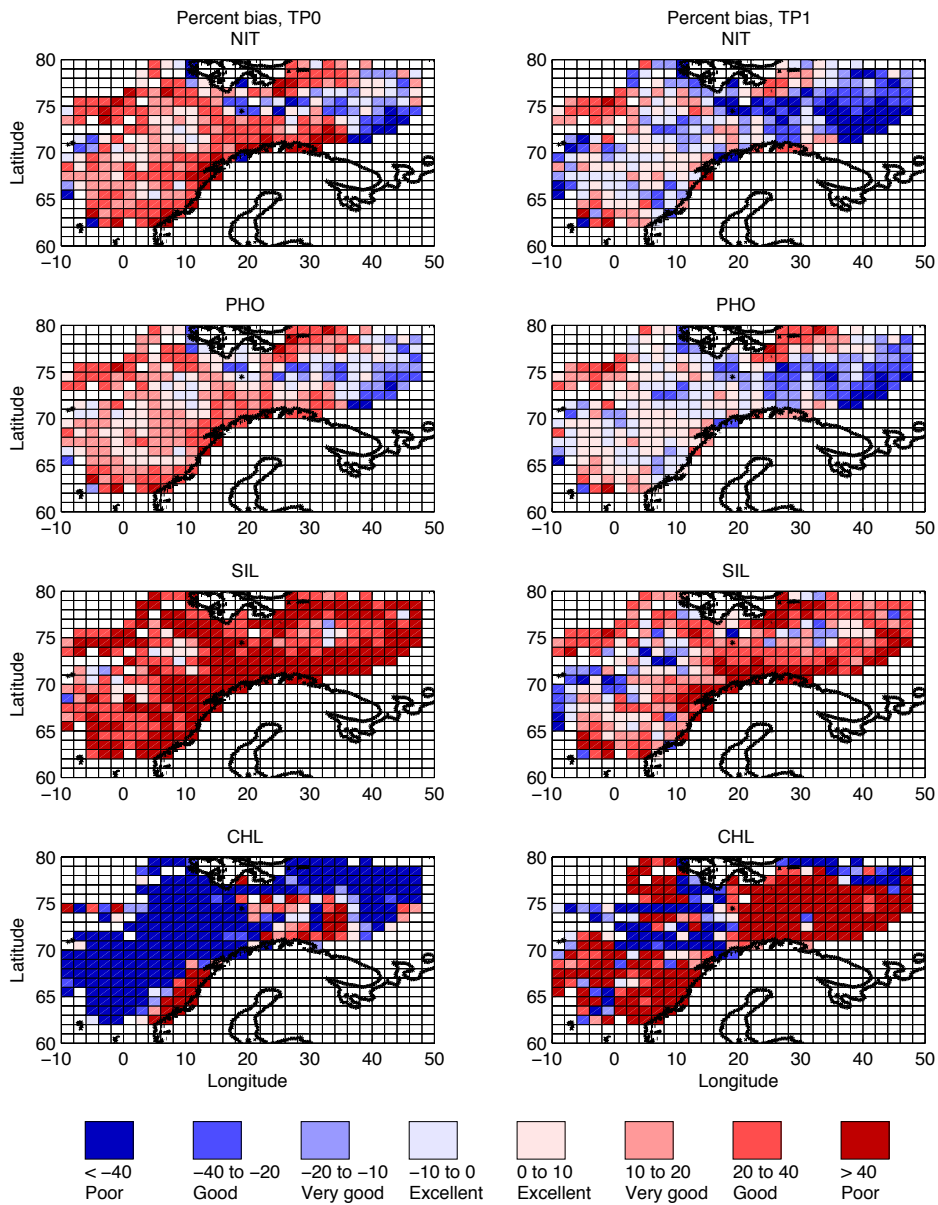
Figure 5. Percentage bias (Pbias, see text) for the model model simulations compared to all available observations from the period 1998-2001.



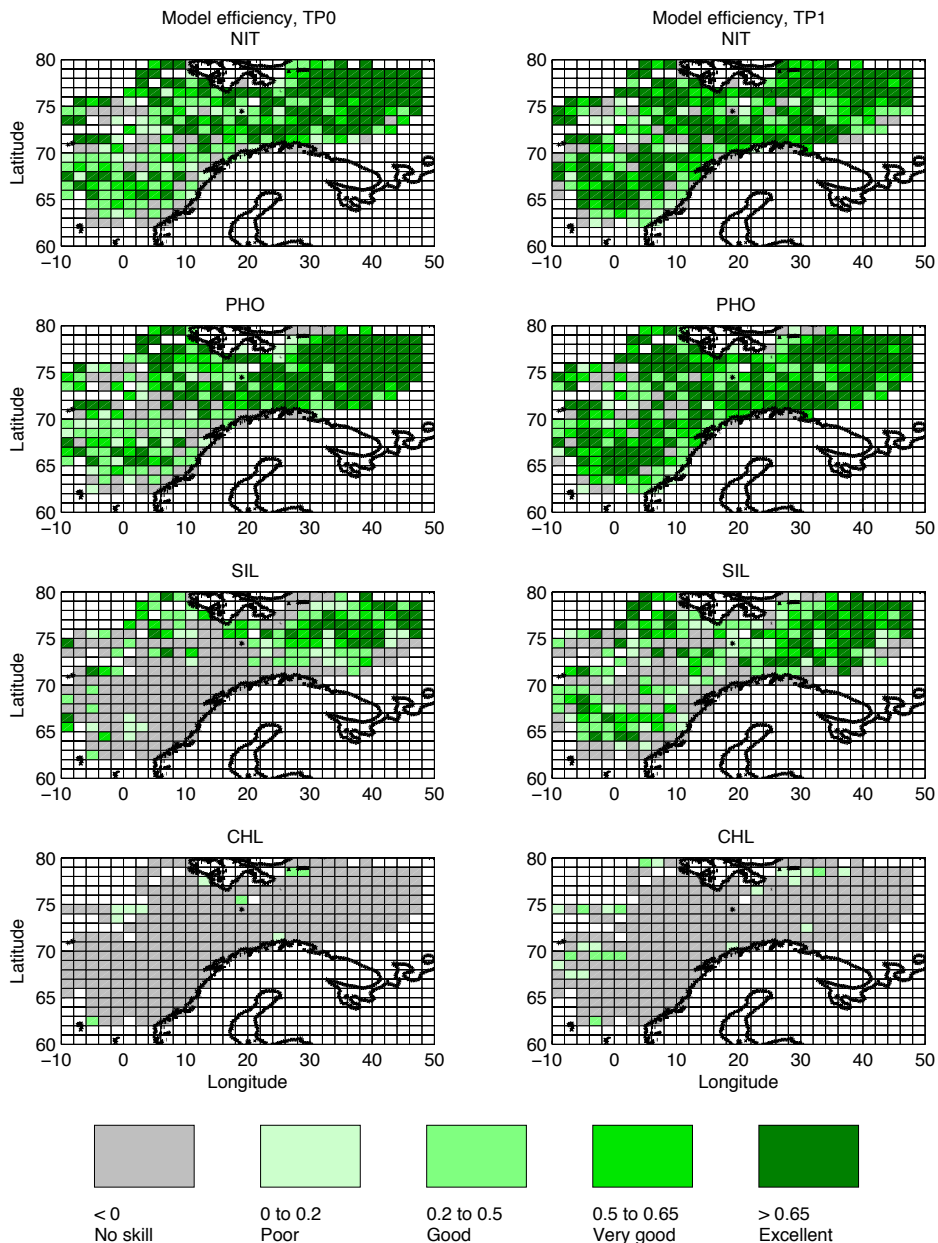
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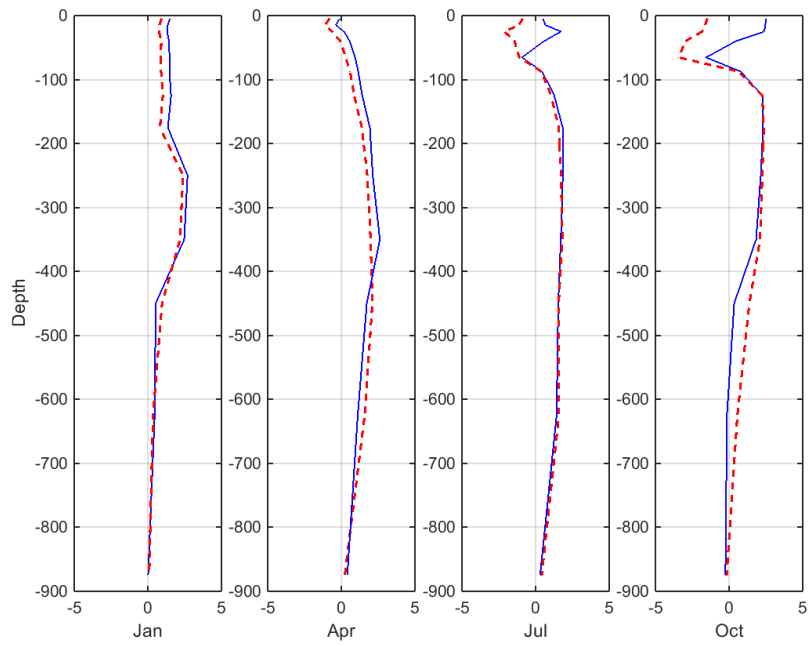
3 Figure 6. Taylor-diagram for comparison with in-situ chlorophyll for the entire area (ALL),  
 4 the Barents Sea (BAS) and the Norwegian Sea including station M (NWS). The curved  
 5 dotted lines show the standard deviation relative to the observations.



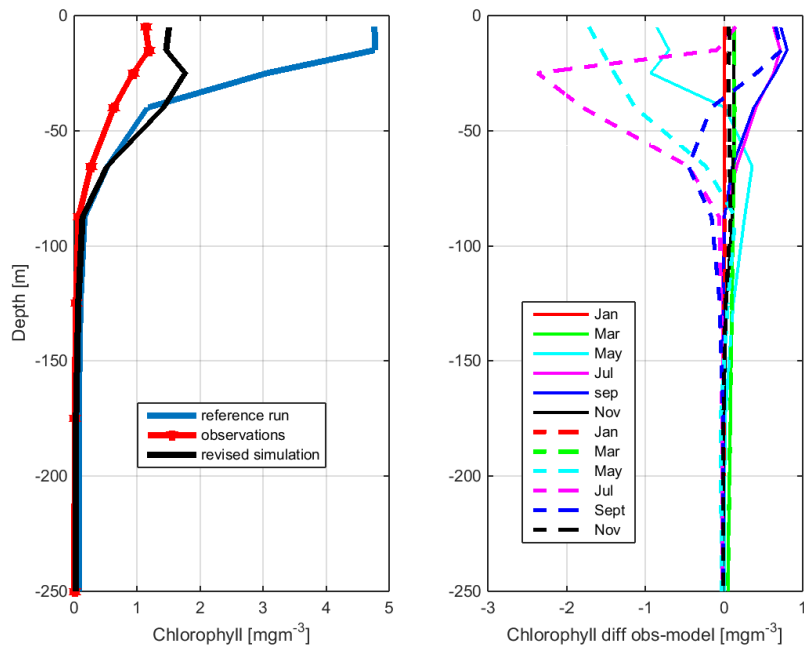
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 2 Figure 7. Percentage bias (Pbias, see text) in the upper 100 meters for the model simulations  
 3 compared to all available observations from the period 1998-2001 in 2x1 degree boxes from  
 4 the simulations with the fine-scale model with the original (TP0) and final set of parameters  
 5 (TP1).



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 2 Figure 8. Model efficiency (ME, see text) in the upper 100 meters for the model simulations  
 3 compared to all available observations from the period 1998-2001 in 2x1 degree boxes from  
 4 the simulations with the fine-scale model with the original (TP0) and final set of parameters  
 5 (TP1).



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 2 Figure 9. Profiles of difference between nitrate [mmol N/m<sup>3</sup>] model and observations in  
 3 different months in the Norwegian Sea box – solid lines (blue) are the revised simulation and  
 4 dashed lines (red) the control run. All observations in the Norwegian Sea box between 1998  
 5 and 2001 have been used.



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2 Figure 10. Chlorophyll profiles from the control and reference run using the higher resolution  
 3 model in June (a) in the Norwegian Sea box as well the difference between observations and  
 4 model in the other months (b) – solid lines are the revised simulation and dashed lines the  
 5 control run. All observations in the Norwegian Sea box between 1998 and 2001 have been  
 6 used.

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**Appendix**

Table A1. Location of changes in the model code, all files are located in  
[https://svn.nersc.no/hycom/browser/HYCOM\\_2.2.12/CodeOnly/src\\_2.2.12/nersc/NORWEC](https://svn.nersc.no/hycom/browser/HYCOM_2.2.12/CodeOnly/src_2.2.12/nersc/NORWEC)  
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	Parameter for tuning	Relevant files	Remarks
N01	Quadratic mortality for phytoplankton	m_NOR05_detritus.F: line 77-89 mod_necessary_ecovars.F90: line 45-54	ZOOPL is 'defined' in all runs in this paper
N02/NO3	Si:N-ratio in diatoms	mod_necessary_ecovars.F90: line 45-54	
N04/NO5/NO6	Meso zooplankton mortality	m_NOR05_zoo_growth.F: line 53	For quadratic mortality, the mortality was set inside the loop calculating mesozooplankton (this code was never submitted to the subversion control system).
N07	Combination of N01 and N02	See above for N01 and N02	
N08/N09/N10	N:Chl-ratio	biocom.h: line 107-108	
N11	Grazing preferences for microzooplankton	m_NOR05_zoo_growth.F: line 26, 100-132	

N12	Grazing preferences for microzooplankton	m_NOR05_zoo_growth.F: line 26, 101	
N13	Combination of N11 and N01	See above for N11 and N01	
N14	Combination of N11 and N2	See above for N11 and N02	
N15	Combination of N14 and N12	See above for N14 and N12	
N16	Combination of N14 and reduced growth rate for phytoplankton	See above for N14 and m_NOR05_affin.F: line 64 and 66	

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