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Reconstructing global Chlorophyll-a variations using a non-linear statistical

- 2 approach
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- 24 **Key Points:**
- A machine learning approach is applied to reconstruct the surface phytoplankton biomass at
- 26 global scale over three decades.
- Chlorophyll variability derived from this statistical approach accurately reproduces satellite
- observations (possibly better than biogeochemical models).

- The sole use of surface predictors allows to accurately reproduce chlorophyll variability, in
- spite of its known sensitivity to three-dimensional processes.

Abstract

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Monitoring the spatio-temporal variations of surface chlorophyll-a concentration (Chl, a proxy of phytoplankton biomass) greatly benefited from the availability of continuous and global ocean color satellite measurements from 1997 onwards. These two decades of satellite observations are however still too short to provide a comprehensive description of Chl variations at decadal to multi-decadal timescales. This paper investigates the ability of a machine learning approach (a non-linear statistical approach based on Support Vector Regression, hereafter SVR) to reconstruct global spatio-temporal Chl variations from selected surface oceanic and atmospheric physical parameters. With a limited training period (13 years), we first demonstrate that Chl variability issued from a 32-years global physical-biogeochemical simulation can generally be skillfully reproduced with an SVR using the model surface variables as input parameters. We then apply the SVR to reconstruct satellite Chl observations using the physical predictors from the above numerical model and show that the Chl reconstructed by this SVR more accurately reproduces some aspects of observed Chl variability and trends compared to the model simulation. The SVR is able to reproduce the main modes of interannual Chl variations depicted by satellite observations in most regions, including El Niño signature in the tropical Pacific and Indian Ocean. In stark contrast with the trends simulated by the biogeochemical model, it also accurately captures spatial patterns of Chl trends estimated by satellites data, with a Chl increase in most extratropical regions and a Chl decrease in the center of the subtropical gyres, although the amplitude of these trends are underestimated by half. Results from our SVR reconstruction over the entire period (1979-2010) also suggest that the Interdecadal Pacific Oscillation drives a significant part of decadal Chl variations in both the tropical Pacific and Indian Ocean. Overall, this study demonstrates that non-linear statistical reconstructions can be complementary tools to in situ and

- satellite observations as well as conventional physical-biogeochemical numerical simulations to
- reconstruct and investigate Chl decadal variability.

1 INTRODUCTION

Phytoplankton—the microalgae that populate the upper lit layers of the ocean—fuels the oceanic food web and regulates oceanic and atmospheric carbon dioxide levels through photosynthetic carbon fixation. The launch of the "Coastal Zone Color Scanner" (CZCS) onboard the Nimbus-7 spacecraft in October 1978 (Hovis et al., 1980) provided the first synoptic view of near-surface chlorophyll-a concentration (Chl, a proxy of phytoplankton biomass). Although primarily focusing on coastal regions, CZCS also provided global pictures of Chl distribution and a new perspective on phytoplankton biomass seasonal variability (Campbell and Aarup, 1992; Longhurst et al., 1995; Yoo and Son, 1998; Banse and English, 2000).

After the failure of CZCS in 1986, ocean color observations were not available for more than a decade. The launch of the modern radiometric Sea-viewing Wide Field-of-View Sensor (SeaWiFS; McClain et al., 2004) in late 1997 followed later by other satellites allowed monitoring and understanding the spatio-temporal Chl variations at global scale over the past two decades. For instance, it revealed that El Niño events induce a Chl decrease in the central and eastern equatorial Pacific in response to reduced upwelled nutrients to the surface layers (e.g., Chavez et al., 1999; Wilson and Adamec, 2001; McClain et al., 2002; Radenac et al., 2012) but also a Chl signature outside the tropical Pacific through atmospheric teleconnections (Behrenfeld et al., 2001; Yoder and Kennelly, 2003; Dandonneau et al., 2004; Messie and Chavez, 2012). It also allowed identifying the Indian Ocean Dipole (IOD; Saji et al., 1999) as the main climate mode driving Chl interannual variations in the Indian Ocean (e.g. Murtugudde et al., 1999; Wiggert et al., 2009; Currie et al., 2013) and monitoring a Chl increase in the subpolar North Atlantic related to the positive phase of the North Atlantic Oscillation (NAO) (Martinez et al., 2016). Aside from the robust Chl decrease monitored in the mid-ocean gyres over the first decade of the XXIst century

(Polovina et al., 2008; Irwin and Oliver, 2009; Vantrepotte and Melin, 2009; Signorini and McClain, 2012), the reliability of the long-term trends derived from these satellite data are more questionable and led to conflicting results in the past literature (Behrenfeld et al., 2006; Vantrepotte and Mélin, 2011; Siegel et al., 2013; Gregg and Rousseaux, 2014). These discrepancies suggest that detection of robust global trend may require several decades of continuous observations (Beaulieu et al. 2013).

The production of longer, consistent ocean color time series can partly alleviate this issue. The combination of the global CZCS and SeaWiFS datasets provided an insight on the Chl response to natural decadal climate variations (Martinez et al., 2009; D'Ortenzio et al., 2012), such as the Pacific Decadal Oscillation (PDO; Mantua et al., 1997) and the Atlantic Multidecadal Oscillation (AMO; Enfield et al., 2001). However, blending these two archives or reconstructing them using compatible algorithms also led to contrasting results (Gregg and Conkright, 2002; Antoine et al., 2005).

The time span of the modern radiometric observations (~20 years), as well as the CZCS-SeaWiFS reprocessed time series, are still too short to investigate Chl decadal variations and longer-term trends. Longer, continuous and consistent records are required. *In situ* biogeochemical observatories can provide such long and continuous records, but their inhomogeneous spatial distribution and varying record length prevent a confident assessment of Chl long-term changes at the scale of a basin (Henson et al., 2016).

Coupled physical-biogeochemical ocean model simulations can provide additional, valuable information's in areas with limited observational coverage. These models resolve reasonably well the seasonal to interannual biogeochemical variability (Dutkiewicz et al., 2001; Wiggert et al., 2006; Aumont et al., 2015). They can however diverge in capturing Chl variations

at a timescale of a decade (Henson et al., 2009a,b; Patara et al., 2011), in particular phytoplankton regime shifts (Henson et al., 2009b). Different biological models are often coupled to different physical models, which renders the attribution of the different modelled responses to their physical or biological components difficult. The decadal or longer variability of the simulated primary producers should then be interpreted cautiously.

In this context, statistical methods reconstructing past Chl variations may be useful alternatives to overcome limitations associated with both observations and numerical models. While statistical reconstructions are now commonly used to extend physical variables back in time (e.g. Smith et al., 2012; Huang et al., 2017; Nidheesh et al., 2017), reconstructions of surface Chl are still in their infancy. Phytoplankton distribution is strongly controlled by physical processes, such as mixing and uplifting, fueling nutrients in the upper-lit layer (i.e., bottom up processes). Thus, relevant physical variables may allow to reconstruct Chl past variations. To our knowledge, a single study allowed the derivation of spatio-temporal surface Chl variations over the 1958-2008 period in the tropical Pacific (Schollaert et al., 2017). This reconstruction used a linear canonical correlation analysis on Sea Surface Temperature (SST) and Sea Surface Height (SSH) to improve the description of the Chl response to the diversity of observed El Niño events and decadal climate variations in the tropical Pacific.

The objective of the present study is to explore the potential of an alternative statistical technique to reconstruct Chl at global scale over a 32-year time-series (i.e., 1979-2010). The considered machine learning technique is based on a Support Vector Regression (SVR) which accounts for non-linearities between predictors and Chl. First, the SVR is trained over 1998-2010 on a self-consistent dataset of physical and Chl variables, all extracted from a forced ocean model simulation that includes a biogeochemical component (i.e., the NEMO-PISCES model). Then,

modelled physical variables are used to reconstruct Chl over 1979-2010. The feasibility and robustness of the proposed reconstruction process is assessed through the comparison of modelled *vs.* reconstructed Chl. In a second step, this framework is applied to satellite ocean color observations.

2 DATA AND METHODS

2.1 The NEMO-PISCES simulation

In this study, we used the "Nucleus for European Modelling of the Ocean" (NEMO) modelling framework (Madec, 2008). The NEMO configuration used displays a coarse resolution with 31 vertical levels and a 2° horizontal grid with a refined 0.5° resolution in the equatorial band. The model includes a biogeochemical component, the Pelagic Interaction Scheme for Carbon and Ecosystem Studies (PISCES; Aumont et al., 2015). PISCES is a model of intermediate complexity designed for global ocean applications (Aumont and Bopp, 2006), which uses 24 prognostic variables and simulates biogeochemical cycles of oxygen, carbon and the main nutrients controlling phytoplankton growth (nitrate, ammonium, phosphate, silicic acid and iron). It simulates the lower trophic levels of marine ecosystems distinguishing four plankton functional types based on size: two phytoplankton groups (small=nanophytoplankton and large=diatoms) and two zooplankton groups (small=microzooplankton and large=mesozooplankton). Chl from PISCES (hereafter referred to as Chl_{PISCES}) is defined as the sum of the simulated diatoms and nanophytoplankton Chl content.

The NEMO-PISCES simulation is forced with atmospheric fields from the interannual Drakkar Forcing Set 5 (DFS5.2, Dussin et al., 2014) for wind, air temperature and humidity, precipitation, shortwave and longwave radiations. It is initialized with the World Ocean Atlas 2005

(WOA05) climatology for temperature, salinity, phosphate, nitrate and silicate (Garcia et al., 2006), while iron initial state is similar to the model climatology employed by Aumont and Bopp (2006). The model simulation was spun up using 3 repetitions of the 30 years' DFS5.2 forcing set, and finally ran over 1979-2010.

Although successfully used in a variety of biogeochemical studies (e.g. Bopp et al., 2005; Gehlen et al., 2006; Lengaigne et al., 2007; Schneider et al., 2008; Steinacher et al., 2010; Tagliabue et al., 2010; Séférian et al., 2013; Aumont et al., 2015; Keerthi et al., 2017; Parvathi et al., 2017 and references therein), the ability of the PISCES model to reproduce satellite surface Chl is briefly illustrated in section 3.1.

2.2 Chl derived from satellite radiometric observations

Satellite surface Chl for Case I waters is provided by the Ocean Colour – Climate Change Initiative (OC-CCI, hereafter referred to as Chloc-cci) from the European Space Agency (http://www.esaoceancolour-cci.org/). This product combines multi-sensor, global, ocean-color products while attempting to reduce inter-sensor biases for climate research (Storm et al., 2013). OC-CCI extends the time series beyond that provided by single satellite sensors and perform better in terms of long-term consistency than other products from multi-mission initiatives (Belo Couto et al., 2016).

Only deep oceanic areas (depth>200 m) are considered to avoid coastal waters where specific non-case-1 waters products are required. The Chl Level-3 product is binned on a regular 1° grid with a monthly resolution over January 1998-December 2010. This time period does not extend beyond 2010 to be consistent with the NEMO-PISCES simulation. Chloc-cci is used to

evaluate the PISCES model performances in Section 3.1, and to train the statistical method in Section 4.

2.3 Predictors and Chl variables

The variability of phytoplankton biomass is driven in many regions of the world ocean and at many timescales by physical processes (e.g., Wilson and Adamec, 2002; Wilson and Coles, 2005; Kahru et al., 2010; Feng et al., 2015; Messie and Chavez, 2015). Our statistical architecture relates to 12 predictors and one biological variable (Chl). A sample thus refers to 13 variables. The 12 predictors (7 physical variables from NEMO-DFS5.2, 2 temporal and 3 spatial parameters) are detailed in **Table 1**, including their influence on Chl variations and the references supporting this influence.

Table 1: physical predictors, the drivers of which they are proxy and associated references.

Proxy used as predictors	Relevance to Chl variations	References			
SST	-Vertical mixing and upwelling	-Behrenfeld et al. (2006); Polovina et al. (2008); Martinez et al. (2009); Thomas et al. (2012)			
	-Impacts on phytoplankton metabolic rates	-Lewandowska et al. (2014)			
Sea Level Anomaly	Thermocline/pycnocline depths	Wilson and Adamec (2001; 2002); Radenac et al. (2012)			
Zonal and meridional wind components	Surface momentum flux forcing and vertical motions driven by Ekman pumping	Martinez et al. (2011); Thomas et al. (2012)			
Zonal and meridional surface current components	Horizontal advective processes	Messie and Chavez (2012); Radenac et al. (2013)			
Short-Wave radiations	Photosynthetically Active Radiation	Sakamoto et al. (2011)			
Month (cos and sin)	Periodicity of the day of the year (day 1 is very similar to day 365 from a seasonal perspective)	Sauzède et al. (2015)			
Longitude (cos and sin)	Periodicity (longitude 0°= longitude 360°)	Sauzède et al. (2015)			
Latitude (sin)					

We purposely limited the predictors to surface variables because our objectives are 1) to reconstruct Chl from physical observations, which are mainly available through remotely sensed surface data (oceanic observations below the surface are indeed usually not accessible at large spatial-scales or interannual time-scales); 2) to build a statistical scheme that can complement more complex numerical models (here, NEMO-PISCES) which simulate complex three-dimensional processes and are costly to run.

First, the SVR is trained on physical predictors from NEMO and DFS5.2 *vs.* Chl_{PISCES}. The reconstructed Chl time-series is referred to as Chl_{Svr-PISCES}. Second, the SVR is trained using the same physical predictors but *vs.* satellite Chl observations (Chloc-cci). The reconstructed Chl time-series is referred to as Chl_{Svr-CCI}.

2.4 Climate indices

Climate indices are provided by the National Oceanic and Atmospheric Administration website (www.esrl.noaa.gov/psd): the AMO, the Multivariate El Niño Southern Oscillation (ENSO) Index (MEI) and the Interdecadal Pacific Oscillation (IPO).

2.5 Support Vector Regression

The statistical reconstruction technique is based on a SVR. This method belongs to kernel methods in Statistical Learning Theory and relates to the Support Vector Machine (SVM, Vapnik, 1998). SVM is a kernel-based supervised learning method (Vapnik, 2000) developed for classification purpose in the early 1990s and then extended for regression by Vapnik (1995). The basic idea behind SVR is to map the variables into a new non-linear space using the kernel function, so that the regression task becomes linear in this space. The learning step estimates the parameters of the regression model according to a linear quadratic optimization problem, which

can be solved efficiently. SVR also uses a robust error norm based on the principle of structural risk minimization, where both the error rates and the model complexity should be minimized simultaneously. Because SVR can efficiently capture complex non-linear relationships, it has been used in a variety of fields, and more specifically for cocanographic, meteorological and climate impact studies (Aguilar-Martinez and Hsieh, 2009; Descloux et al., 2012; Elbisy, 2015; Neetu et al., 2020), as well as in marine bio-optics (Kim et al., 2014; Hu et al., 2018; Tang et al., 2019).

Predictors and Chl are normalized by removing their respective average and dividing them by their standard deviations. Two SVR are trained over 1998-2010: one on Chlpisces and one on Chloc-cci (Step A in Figure 1). This time period has been chosen as 1998 is the first complete year of the satellite Chloc-cci time-series, and 2010 is the last year available of the modelled Chlpisces. The two resulting SVR schemes are applied on the NEMO-DFS5.2 physical predictors over 1979-2010. Finally, the annual means and standard deviations initially removed are applied to perform the back transformation and reconstruct either Chlsvr-Pisces or Chlsvr-Cci (Step B in Figure 1).

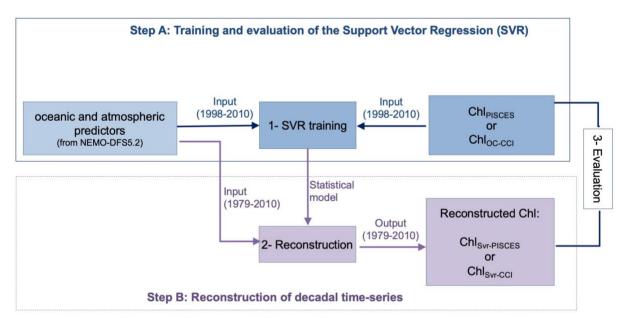


Figure 1 | Steps performed to train the SVR and reconstruct Chl time-series

Considering a Gaussian kernel, SVR only involves the selection of two hyperparameters: the penalty parameter C of the error term and the kernel coefficient gamma, driving the reduction of the cost function. C and gamma values are 1 and 0.1, respectively when the SVR is trained on Chlpisces, and 2 and 0.3 when trained on Chloc-cci (See details in the Supplementary material, and Figure Supp 1A). Sensitivity tests to an increasing portion of the sample total number (from 0.2 % to 9 % of the full dataset) used in the training process are performed (See Supplementary materials and Figure Supp 1B). The mean absolute error stabilizes for a sample number higher than 6 %, suggesting that the SVR skills don't improve much afterward. This observation combined with computational limitations lead us to present the 9% experiment hereafter.

2.6 Empirical Orthogonal Function analysis

The SVR skills to reconstruct Chl interannual to decadal variations are investigated performing Empirical Orthogonal Function analysis on Chlpisces, Chloc-cci, Chlsvr-pisces and Chlsvr-cci. First, Chl data are centered and reduced (i.e., the monthly climatology is removed and the induced anomalies are divided by their standard deviations) to avoid an overly dominant contribution of high values on the analysis (Emery and Thompson, 1997) over the periods of interest (i.e., 1998-2010 or 1979-2010). A 5-month running mean is applied to focus on the interannual/decadal signal. The analysis is separately performed for the Atlantic, Pacific and Indian Oceans north of 40°S until 60°N, and for the 40°S-60°S region hereafter referred to as the Austral Ocean. Indeed, the large area covered by the Pacific Ocean and its dominant modes in climate variability (i.e., ENSO/IPO), could regionally dampen other modes of variability. Basin-scale spatial maps are then gathered to a global one, refer to as EOF. The associated time-series refer to as the Principal Components (PCs).

3 Synthetic reconstruction from a physical-biogeochemical ocean

model

This section assesses the reliability and robustness of the SVR approach using a complete and coherent dataset extracted from a global simulation performed with a coupled physical-biogeochemical ocean model. The SVR is first trained over 1998-2010 on Chlpisces, and Chlsvr-Pisces is reconstructed over 1979-2010. Chlpisces and Chlsvr-Pisces are then compared over 32 years to evaluate the consistency of the proposed data-driven reconstruction scheme.

3.1 Evaluation of Chlpisces at global scale

The ability of the NEMO-PISCES model to reproduce the satellite Chl over 1998-2010 is briefly presented here. Boreal winter and summer climatology from Chlpisces compare reasonably well with those of Chloc-cci (Figures 2A vs. B, and C vs. D). The model correctly represents the main spatial patterns with, for instance, higher Chl and a stronger seasonal cycle at high latitudes, despite an overestimated biomass in the Southern Ocean (Launois et al., 2015). The model also captures low Chl in the subtropical gyres, with some underestimation. This discrepancy may be explained by the lack of acclimation dynamics to oligotrophic conditions or by the assumption of constant stoichiometry either in phytoplankton or in organic matter in the model (Aumont et al., 2015; Ayata et al., 2013). The model underestimates Chl values in the equatorial Atlantic and Arabian Sea. In this latter region, mesoscale and submesoscale processes unresolved by the model have been shown to be of critical importance (Hood et al., 2003; Resplandy et al., 2011). Finally, the parameterization of nitrogen-fixing organisms not explicitly modelled in that PISCES version could explain the Chlpisces underestimation in the western Pacific in austral summer (Dutheil et al., 2018).

High Chl are accurately simulated in the eastern boundary upwelling systems. In two of the three main High Nutrient Low Chlorophyll (HNLC) regions, i.e. the equatorial Pacific and the eastern subarctic Pacific, the model successfully reproduces the moderate Chloc-cci. However, the model overestimates Chloc-cci east of Japan because of an incorrect representation of the Kuroshio current trajectory. This common bias in coarse resolution models (i.e., Gnanadesikan et al., 2002; Dutkiewicz et al., 2005; Aumont and Bopp, 2006) is potentially related to too deep mixed layer simulated in winter inducing very strong spring blooms (Aumont et al., 2015). In the Southern Ocean, the third and largest main HNLC region, the model overestimates Chloc-cci values, especially during summer. However, the standard satellite algorithms that deduce Chl from reflectance tend to underestimate *in situ* observations by a factor of about 2 to 2.5, especially for intermediate concentrations (e.g., Dierssen and Smith, 2000; Kahru and Mitchell, 2010). It is to note that Chl in physical-biogeochemical coupled models is commonly overestimated in the Southern Ocean, and systematically underestimated in the oligotrophic gyres (Séférian et al., 2013).

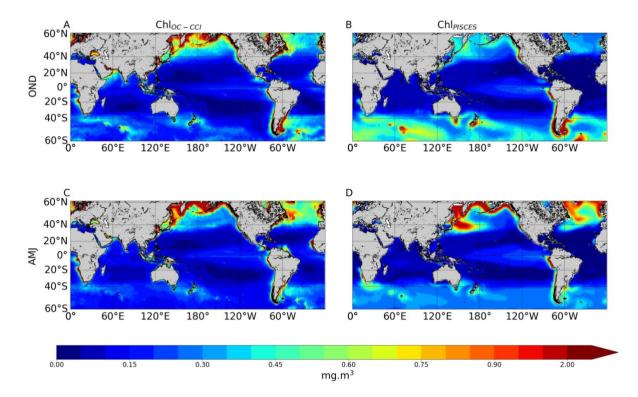


Figure 2 | Surface seasonal mean of Chl (mg.m⁻³) over 1998-2010 derived from satellite (left panels) and the PISCES model (right panels), in October-November-December (**A-B**) and April-May-June (**C-D**).

The 1st mode of the EOF analysis performed on interannual Chl displays close percent of total variance for Chloc-cci and Chlpisces (16.6% vs. 21.1%, respectively). Their PCs in the Pacific Ocean are well correlated with the MEI (r = 0.71 and 0.89 with p=0.0015 and p<0.001, respectively; **Figure 3C**). PCs show the greatest positive values in January 1998 during the peak of the strong 1997/1998 El Niño event and the greatest negative values during the following La Niña beginning of 1999. The associated EOFs display a Chl horseshoe pattern (**Figure 3A and 3B**), reminiscent of the ENSO pattern on SST from NEMO (SST_{NEMO}; **Figure Supp 2**; Messie and Chavez, 2012). While the tropical Pacific experiences a Chl decrease during El Niño events, the North and South Pacific display a Chl increase, and inversely during La Niña. This typical ENSO pattern is also related to remote Chl anomalies outside the Pacific induced by atmospheric teleconnections, such as a Chl decrease in the tropical North Atlantic and in the South Indian Ocean

during El Niño. Although the Atlantic and Indian Ocean's PCs are not correlated with the MEI (0.14 and 0.05, respectively), their EOFs are similar to those obtained from analysis performed at global scale (*vs.* basin scale here) and which have been largely discussed in the past (e.g., Behrenfeld et al., 2001, 2006; Chavez et al., 2011; Yoder and Kennelly, 2003). Chlpisces reasonably well captures the first mode of Chloc-cci interannual variability over 1998-2010 in the Pacific and Atlantic Oceans, with 0.89 and 0.77 (p<0.001) correlations between their PCs, respectively, but not in the Indian Ocean, where the PCs correlation is far weaker (0.13) and insignificant (**Figure 3C-E**).

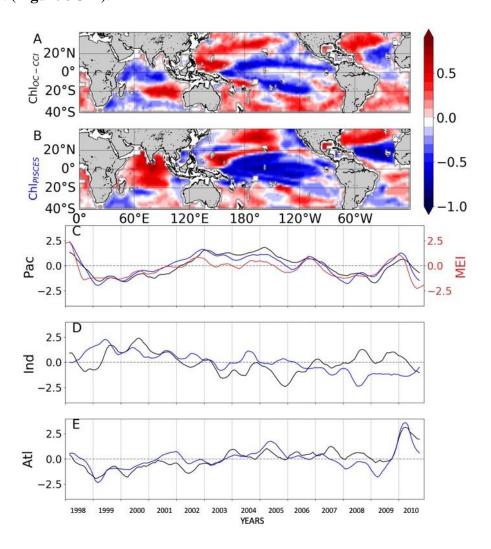


Figure 3 | 1^{st} mode of basin-scale EOFs of interannual **A**) Chloc-cci and **B**) Chl_{PISCES}, and their corresponding PCs over 1998-2010 in the **C**) Pacific, **D**) Indian and **E**) Atlantic Oceans. Chloc-cci and Chl_{PISCES} PCs are represented by the black and blue lines, respectively. The MEI index is reported in red (right y-axis) on C.

3.2 Evaluation of the SVR method trained on synthetic data only

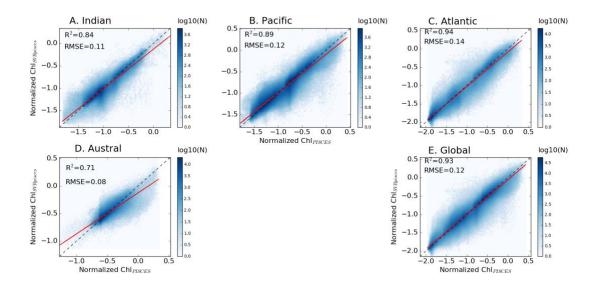
3.2.1 Statistical performances

A first evaluation of the SVR applied on the synthetic dataset (i.e., both physical and biogeochemical model outputs) is provided for the dedicated subset (i.e., 20% of 9% of the total data set) over the 1998-2010 training time period. Chlpisces and Chlsvr-Pisces datasets display a determination coefficient of 0.95 and a root mean square error (RMSE) of 0.22 (see **Figure Supp 1C**), indicating at first glance a very good ability of the SVR to reconstruct Chlpisces. The SVR reconstruction is very accurate when comparing the full modelled and reconstructed Chl for (i) the 1998-2010 training time period, (ii) the 1979-1997 fully independent dataset and (iii) the 1979-2010 whole dataset, both at global and basin scales (**Table 2** and **Figure 4**). For each oceanic basin, determination coefficients between both datasets exceed 0.85, except in the Austral Ocean where they get down to 0.71. RMSE are lower than 0.14 and associated with a slope ranging from 0.84 in the Austral to 0.97 in the Atlantic (**Figure 4**). In addition, the quality of the reconstructed Chlsvr-Pisces over the 1979-1997 independent time period is only marginally degraded compared to the 1998-2010 training period or the 1979-2010 full period.

Table 2 | Statistical performances between Chlpisces vs. Chlsvr-Pisces normalized monthly anomalies for the global ocean and the 4 oceanic basins over the 1998-2010, 1979-1997 and the whole 1979-2010 time period. The determination coefficient, NRMSE and the slope of the Chl_{PISCES} vs. Chl_{Svr-PISCES} regression line are indicated.

	1998-2010				1979-1997				1979-2010			
	R ²	RMSE	slope	number of bins	R ²	RMSE	slope	number of bins	R ²	RMSE	slope	number of bins
Global 60°S-60°N	0.96	0.1	0.95	4 487 457	0.94	0.12	0.95	6 562 272	0.94	0.11	0.95	11 049 729
Pacific	0.92	0.1	0.93	1 945 173	0.88	0.12	0.9	2 843 501	0.9	0.11	0.91	4 788 674
Indian	0.86	0.11	0.85	612 612	0.85	0.11	0.89	895 356	0.85	0.11	0.87	1 507 968
Austral	0.81	0.06	0.8	1 026 876	0.71	0.08	0.75	1 506 260	0.76	0.07	0.77	2 533 136
Atlantic	0.96	0.12	0.96	902 796	0.94	0.14	0.94	1 317 155	0.94	0.13	0.95	2 219 951





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Figure 4 | Scatter plots of Chlpisces vs. Chlsvi-Pisces normalized monthly anomalies over 1979-1997, A-D) for each basin and E) at global scale between 60°S and 60°N. The Chlpisces vs. Chlsvr-Pisces and the 1:1 regression lines are plotted as the red and black lines, respectively. The figure is color-coded according to the density of observations.

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Evaluation of the reconstructed Chl spatio-temporal variability

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The Normalized Root-Mean-Square-Error (NRMSE, i.e. RMSE normalized by the average of Chl used to train the SVR) between Chlpisces and Chlsvr-Pisces filtered with a 5-month running

mean (to discard the high frequency signal) shows an error ranging between 10 and 20% over 1998-2010 (**Figure 5A**). Their correlation exceeds 0.7 (p<0.001) over most of the global ocean (**Figure 5B**). At mid-latitudes they are generally larger than 0.8, and they range between 0.6 and 0.9 in the equatorial Pacific. This accurate reconstruction demonstrates that a strong relationship exists between physical processes and Chl at global scale. However, the reconstructed Chl field can be regionally less accurate. For instance, the edges of the oligotrophic gyres (delimited by the 0.1 mg.m⁻³ contour in **Figure 5A**) exhibit the highest NRMSE and lowest correlations. Large NRMSE are also evident in the Gulf Stream region while the western tropical Atlantic exhibits lower correlations than 0.5.

Those discrepancies could be due first to the zooplankton grazing pressure (top-down control) which is often overestimated in PISCES simulations. It results in an underestimated nanophytoplankton biomass in the oligotrophic gyres, emphasized along their edges (Laufkötter et al., 2015). Because the top-down control is not accounted for by the SVR, Chl variability induced by the overgrazing in these areas might not be captured. Second, in the equatorial Pacific Ocean, a minimum iron threshold value has been imposed (0.01 nmol.L⁻¹) in the biogeochemical model. Without that threshold Chl is too low on both sides of the equator, resulting in a strong accumulation of macronutrients and a spurious poleward migration of the subtropical gyre boundaries (Aumont et al., 2015). While the existence of such a threshold suggests that a minor but regionally important source of iron is missing in PISCES, it also suggests the inability of the SVR in reproducing ecosystem dynamics related to such artificial input of micro-nutrient. Finally, atmospheric input of iron through desert dust deposition is known to be stronger in the Atlantic than in the Pacific Ocean (Jickells et al., 2005). Such signal cannot be accounted for by the SVR

with the given predictors, which might (with meso – and sub-mesoscale activities) explain the higher NRMSE in the north western Atlantic than in the north-western Pacific.

As expected, areas of high NRMSE and low correlations between Chl_{PISCES} and Chl_{Svr-PISCES} identified over 1998-2010 (**Figure 5, left column**) extend and strengthen over 1979-1997 (**Figure 5, right column**). Indeed, the correlations significantly decrease in the tropical Pacific while they slightly decrease in mid-latitudes between the two periods. Correlations remain high and NRMSE low in the North-West Pacific, North and South-West Atlantic, and South Indian Oceans as well as over a large part of the Southern Ocean providing confidence for analyses extended beyond the training period of the SVR.

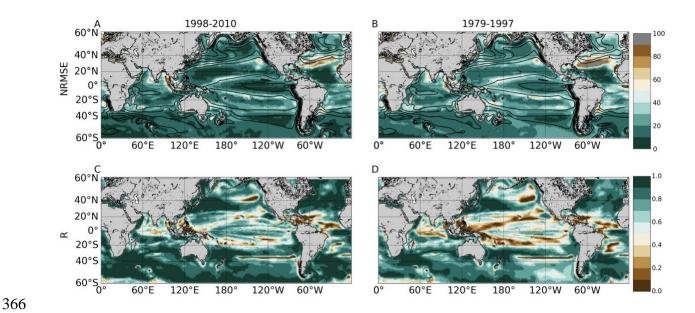


Figure 5 | **A-B**) NRMSE (in %) and **C-D**) correlation between Chl_{PISCES} *vs.* Chl_{Svr-PISCES} after applying a 5 month-running mean on both time-series. These 2 diagnostics are calculated over 1998-2010 (left column) and 1979-1997 (right column). Contours on the upper panels show their respective 1998-2010 Chl time average (every 0.1 mg.m⁻³).

The analysis is now extended to the 1979-2010 time-period to investigate the skills of the SVR in reproducing phytoplankton interannual/decadal cycles. The 1st EOFs of Chlpisces *vs*.

Chlsvr-PISCES have the same sign of variability over 72% of the global ocean (**Figures 6A-B**). Both EOFs are similar in the Pacific and Atlantic Oceans and their PCs are highly correlated over 1979-2010 (**Table 3; Figures 6C,6E**). In the Pacific, these EOFs strongly resemble the typical horseshoe pattern of IPO with SST anomalies of opposite polarities in the tropical and extra-tropical Pacific regions (**Figure Supp 3**). Correlations between ChlPISCES and Chlsvr-PISCES 1st PCs and the IPO index are high (0.94 and 0.95 with p<0.001, respectively; blue and black *vs.* red lines in **Figure 6C**). It highlights that the 1st mode of Chl variability in the Pacific is strongly driven by the IPO. In the Atlantic, both PCs are strongly correlated with the AMO (-0.8 for Chlsvr-PISCES and -0.85 for ChlPISCES with p<0.001; **Figure 6E**). The AMO shifts from a cold to a warm phase in the mid-1990's (**Figure Supp3**), and is associated with a decrease in Chl (**Figures 6A-B**).

The 1st two modes explain a similar percent variance for Chl_{PISCES} and Chl_{Svr-PISCES} in the four oceanic basins, with the exception of the 1st mode in the Atlantic Ocean (see **Table 3**). In this basin Chl_{Svr-PISCES} percent variance is underestimated by a factor 2 compared to Chl_{PISCES}, while their 1st EOFs and PCs are well correlated. One explanation might be that the AMO is the climate cycle with the longest period (80 years) when compared to the IPO. Thus, it might be the most difficult signal to reproduce as the SVR is trained over a relatively "short" 12 years' time-period.

The agreement between Chlpisces and Chlsvr-Pisces 1st EOFs is not as good in the Austral and Indian Oceans when compared to the Atlantic and Pacific Oceans (**Table 3** and **Figures 6A-B**). In the Indian Ocean, the Chlpisces EOF exhibits a maximum positive variability along the western Arabian Sea, while it is located north-east of Madagascar for Chlsvr-Pisces. In the Austral Ocean, Chlpisces and Chlsvr-Pisces EOFs roughly follow a zonal distribution.

A strong correspondence between SST and Chl has been previously reported over a large part of the global ocean (Behrenfeld et al., 2006; Martinez et al., 2009; Siegel et al., 2013),

demonstrating the close interrelationship between ocean biology and climate variations. Consequently, it is not surprising to observe strong correlations between Chl_{PISCES} or Chl_{Svr-PISCES} and climatic indexes mostly built on SST anomalies (**Figure Supp 3**).

Table 3 | Percent variance explained by the first two modes of the Empirical Orthogonal Function analysis performed on Chl_{PISCES} and Chl_{Svr-PISCES} for each oceanic basin over 1979-2010. The correlation (r) between the Chl_{Svr-PISCES} and Chl_{PISCES} PCs is also reported with a significant level of p<0.001 (*) and p<0.002 (**).

2nd mode

	Chl _{PISCES}	Chl _{Svr-PISCES}	r	Chl _{PISCES}	Chl _{Svr-PISCES}	r
Pacific	19.7	23.5	0.95*	7.8	6.7	0.6*
Indian	13.1	14.1	0.58*	10.5	9.4	0.78*
Austral	13.8	12.1	0.62*	11	8.4	0.47**
Atlantic	23.2	13.9	0.81*	9.6	10.9	0.73*

1st mode

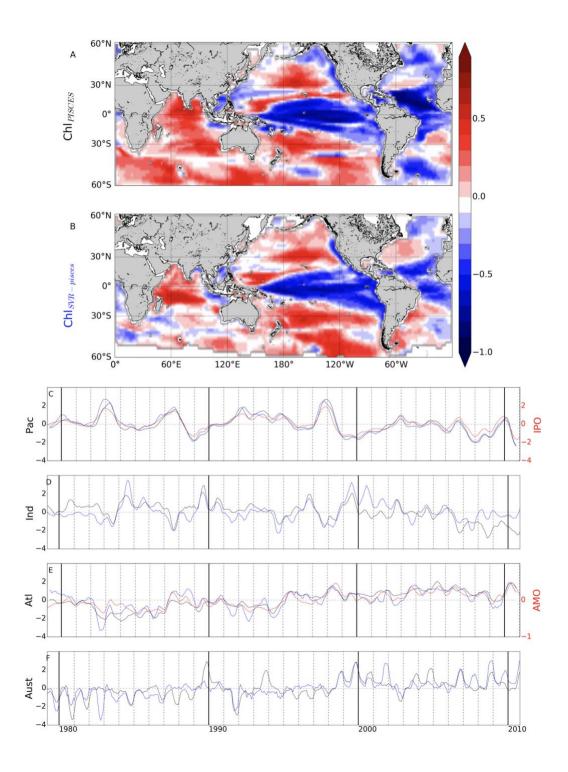


Figure 6 | 1^{st} mode of basin-scale EOFs of interannual **A**) Chl_{PISCES} and **B**) Chl_{Svr-PISCES}, and their corresponding PCs over 1979-2010 in the **C**) Pacific, **D**) Indian **E**) Atlantic and **F**) Austral Oceans (black and blue lines, respectively). Climate indices are reported in red (right y-axis).

The 2nd mode of variability of Chl_{PISCES} is also well reproduced by the SVR. The percent variances are close (**Table 3**) as well as their spatio-temporal variability in the four oceanic basins (**Figure Supp 4**). The high correlations between the first two modes of Chl_{PISCES} vs. Chl_{Svr-PISCES} highlight the SVR ability to relatively well reproduce the Chl_{PISCES} low-frequency variability.

4 Application to satellite radiometric observations

4.1 SVR statistical performances and sensitivity tests

In this section, the SVR uses the same physical predictors from NEMO-DFS5.2 as in Section 3, but it is trained on satellite radiometric observations (e.g., Chl_{OC-CCI}). The same procedure is followed (See **Figure Supp 5A-B**). A first validation is performed for 20% of 9% of the full data set and over the 1998-2010 training period showing a high determination coefficient of 0.87 and RMSE of 0.37 between Chl_{OC-CCI} and Chl_{Svr-CCI} (**Figure Supp 5C**).

As expected, the regression lines of Chloc-cci vs. Chlsvr-cci for each oceanic basin and at global scale are farther away from the 1:1 line than for the synthetic study over the training period, but still remain close (higher slope than 0.8, except in the Austral Ocean; **Figure 7**). The SVR trained on NEMO-DFS5.2 predictors vs. satellite Chl is expected to be less efficient than the SVR trained on the coherent NEMO-DFS5.2-PISCES physical-biogeochemical dataset. Some of the biological interactions/processes (such as the diversity of the prey-predator relationships, the complexity of photoacclimation phenomena) are not yet optimally formulated by model equations inducing that Chl derived from numerical modelling is oversimplified compared to the complexity of the real ocean. Not to mention that satellite Chl may itself be partially affected by other components that are not Chl, such as colored dissolved organic matter (CDOM; Morel and Gentili, 2009) and suspended particulate matter (SPM). Phytoplankton can also adjust their intracellular

Chl according to light and nutrient availability (e.g., Laws and Bannister, 1980; Behrenfeld et al., 2015). The induced Chl changes are no longer ascribed to changes in biomass. All these signatures on satellite Chl could explain Chl_{Svr-CCI} underestimation. Nevertheless, correlations between Chl_{Svr-CCI} and Chl_{OC-CCI} remain high over the training time period (0.92, 0.94 and 0.93 for the Indian, Pacific and Atlantic Oceans, respectively, **Figure 7**).

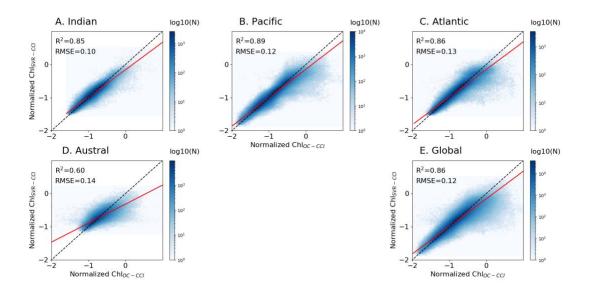


Figure 7 | Same as Figure 4 for Chloc-cci vs. Chl_{Svr-CCi} normalized monthly anomalies over 1998-2010.

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The NRMSE between Chloc-cci vs. Chl_{Svr-cci} is lower than 20% over most of the global ocean (**Figure 8A**). Correlations higher than 0.9 (p<0.001) are evident over large subtropical areas in the Atlantic, Indian and Pacific Oceans as well as in the Equatorial Pacific (**Figure 8B**). Interestingly, the SVR generally does a better job at reconstructing the satellite Chl than the modeled one (**Figure 5A-B** vs. **Figure 8**). NRMSE are higher at high latitudes and along the oligotrophic area boundaries, although to a less extent than for Chl_{PISCES}. Because Chloc-cci can only be retrieved under clear sky conditions, gaps in satellite observations (especially during

wintertime) likely alters the SVR learning and could explain such a degradation of Chl_{Svr-CCI} as moving towards high latitudes.

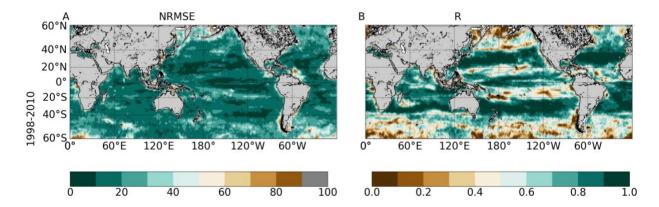


Figure 8 | **A**) NRMSE (in %) and **B**) correlation between Chloc-cci vs. Chlsvr-cci over 1998-2010 after applying a 5 month-running mean on both time-series. Contours on the NRMSE show the 1998-2010 Chloc-cci time average (every 0.1 mg.m⁻³). Correlations <0.73 and 0.6 are significant with a p-value<0.001 and 0.01, respectively.

4.2 Reconstruction of satellite Chl interannual to decadal variability and trends

The SVR ability to replicate Chloc-cci interannual variability is now investigated over 1998-2010 (**Figure 9**). In the Pacific Ocean, Chloc-cci and Chlsvr-cci 1st EOFs are close (**Figure 9A** *vs.* **9B**), their PCs are highly correlated (r=0.89, p<0.001; **Figure 9C**), and their percent variance are similar (**Table 4**). As presented in Section 3.1, this mode of Chl variability can be attributed to ENSO, given their EOFs pattern as well as their PCs highly correlated with the MEI (roc-cci/mei=0.71 and rsvr-cci/mei=0.91, with p=0.0015 and p<0.001, respectively). Interestingly, Chlsvr-cci EOFs are closer to Chloc-cci than Chlpisces in several areas such as in the north-western Pacific, the south-western Atlantic and the Indian Ocean from Madagascar to the western coast of Australia (**Figures 9A-B** *vs.* **Figure 3B**). Consistently, correlations between Chloc-cci and Chlsvr-cci PCs in the three basins and for the 1st two modes are higher than between Chloc-cci and Chlpisces (**Table 4**).

Table 4 | Percent variance explained by the first two modes of the Empirical Orthogonal Function analysis performed on Chloc-cci, Chlsvr-cci and Chlpisces for each oceanic basin over 1998-2010. The correlation (r) between Chloc-cci vs. Chlsvr-cci and between Chloc-cci vs. Chlpisces is also reported with a significant level of P<0.001 (*) and P<0.02 (**).

			1st mode			2 nd mode					
	% of variance			r Chl oc.ccivs.		% of variance			r Chl occes vs.		
	Chloc-cci	Chl _{Syr-CCI}	Chlpisces	Chl _{syr-CCI}	Chlpisces	Chloc-cci	Chl _{svr-CCI}	Chlpisces	Chl _{syr-CCI}	Chlpisces	
Pacific	16.6	23.7	21.1	0.89*	0.89*	10.7	12.5	13.6	0.81*	0.52**	
Indian	16.9	16.6	17.3	0.57**	0.13	11.8	12.2	15.1	0.48	0.36	
Atlantic	14	17.9	19.4	0.85*	0.77*	10.7	9.1	12.5	0.82*	0.59**	

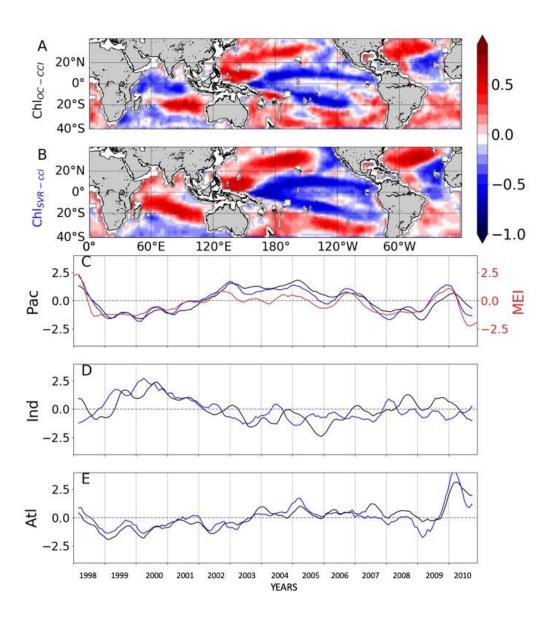


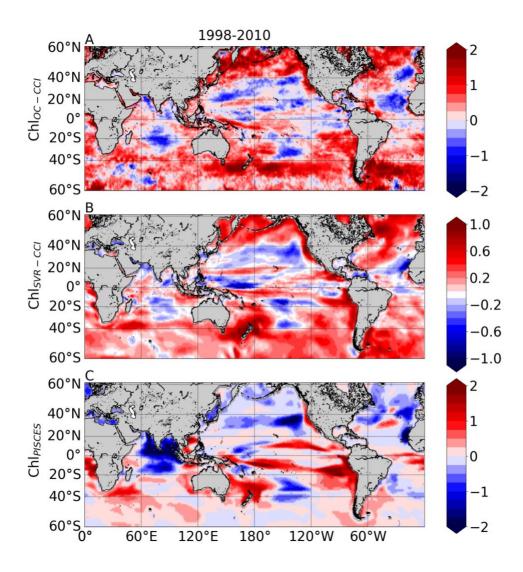
Figure 9: 1st mode of basin-scale EOFs of interannual **A**) Chloc-cci and **B**) Chl_{svr-cci} and their associated PCs over 1998-2010 in the **C**) Pacific, **D**) Indian and **E**) Atlantic Oceans as the black and blue lines, respectively (left y-axis). The climate indices are reported in red on the right y-axis.

Chloc-cci linear trends over 1998-2010 exhibit large areas of increase or decrease (red and blue areas in **Figure 10A**, respectively). Productive regions at high latitudes and along the equatorial and upwelling areas generally exhibit positive Chloc-cci trends, albeit many underlying regional nuances. Contrastingly, trends are generally negative in the center of the gyres. These regional trends are consistent with those extracted from the first 13 years of the SeaWiFS record and discussed by Siegel et al. (2013) (see their Figures 5b and 8b). The negative trends in the oligotrophic gyres were also reported by Signorini et al. (2015) who attributed this behavior to MLD shallowing trends. Surface water density variability induced by changes in temperature and salinity, combined with wind stirring, are effective drivers of vertical mixing, which in turn control the renewal of nutrients from the rich-deep layers toward the euphotic zone. Thus, shallower MLD would decrease nutrient uplift and phytoplankton growth in the oligotrophic areas.

Chlsvr-cci trends agree qualitatively well with those of Chloc-cci at global scale (**Figures 10B** *vs.* **10A**, respectively). Indeed, decline of Chlsvr-cci can be observed in the center of the gyres, while outside Chlsvr-cci generally increases in a similar way to Chloc-cci. Chlsvr-cci accurately captures the largest Chloc-cci increase observed in the Southern Ocean along the Antarctic Circumpolar Current. While Gregg and Casey (2004) reported a substantial negative bias in the SeaWiFS data for this region when compared to *in situ* observations, which could hamper the reliability of satellite trends discussed in this area (e.g. Siegel et al., 2013), the SVR remains able to reproduce the positive observed trend. Despite qualitative spatial agreements, it is noteworthy

that the SVR underestimates by half the magnitude of the satellite trend (see scales in **Figure 10A** *vs.* **10B**).

Interestingly, trends in Chlpisces generally differs from Chloc-cci (**Figure 10C**). This is striking for the North Pacific and Atlantic high latitudes, but also in the equatorial Atlantic and Arabian Sea with opposite trends when compared with Chloc-cci and Chlsvr-cci, and in a more mitigated manner in the Austral Ocean.



 $\label{eq:Figure 10} \textbf{Figure 10} \mid Linear \ trends \ (in \ \% \ year^{-1} \) \ calculated \ over 1998-2010 \ from \ the \ monthly \ \textbf{A}) \ ln(Chloc-cci), \ \textbf{B}) \ ln(Chl_{Svr-CcI}), \ \textbf{C}) \ ln(Chl_{PISCES}). \ Note that \ the \ scale \ is \ divided \ by \ 2 \ for \ ln(Chl_{Svr-CcI}).$

Chlsvr-CCI is also compared with the only historical consistent dataset built by Antoine et al. (2005) who reanalyzed ocean color time series from CZCS (1979–1983) and SeaWiFS (1998–2002). A 22% global mean increase of Chl between the two era was reported. It was mainly due to large increases in the intertropical areas and to a lesser extent in higher latitudes, while oligotrophic gyres displayed declining concentrations (**Figure 11A**). SST from the SODA reanalysis was used as a proxy of ocean stratification and opposite Chl and SST changes over 60% of the ocean between 50°S and 50°N was reported (light blue and yellow in **Figure 11B**, adapted from Martinez et al., 2009). This inverse relationship was used to hypothesized that multidecadal changes in global phytoplankton abundances were related to basin-scale oscillations of the ocean dynamics. Briefly, SST changes were related to a regime shift of the PDO (although the use of the basin-scale IPO would have been more appropriate) from a warm to a cold phase in the Pacific and Indian Oceans leading to an increase of Chl, and inversely in the Atlantic Ocean with a regime shift from a cold to a warm phase of the AMO leading to a Chl decrease.

Observed Chl changes over the last decades are accurately reproduced by Chlsvr-CCI, including a Chl increase in the equatorial Pacific and the southern tropical Indian Oceans, as well as a Chl decline in both the Atlantic and Pacific oligotrophic gyres (**Figure 11C**). However, the magnitude of the SVR reconstructed Chl is underestimated (note that the Chl ratio is multiplied by 2 in **Figure 11C** to allow the comparison with **Figure 11A**). On average, the inverse relationship between Chlsvr-CCI and SSTNEMO (**Figure Supp 4**) occurs over 69.4% of the global ocean between 50°S and 50°N in a similar way to that reported by Martinez et al. (2009), especially in the Pacific Ocean (see **Figure 11D** *vs.* **Figure 11B**). In the Indian Ocean, although Chl mainly increases in both studies, it is here associated with a SST decrease. Interestingly, this inverse Chl-SST relationship in the Indian Ocean (yellow area in **Figure 11D**) was reported in Behrenfeld et al.

(2006) over the SeaWiFS era, suggesting that the SST dataset used in Martinez et al. (2009) may have decadal discrepancies for this region.

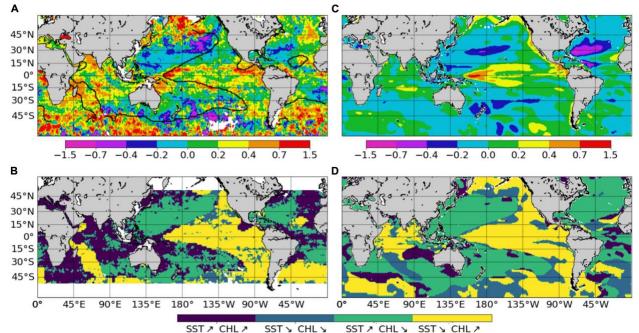


Figure 11 | Chl change from the CZCS (1979–1983) to the SeaWiFS (1998–2002) era, expressed as the logarithm of the ratio of the average values over the two time periods **A**) from satellite Chl adapted from Antoine et al. (2005), **C**) from Chl_{Svr-CCI}. Note that this ratio is multiplied by 2 to fit the same color bar as in **A**. Maps of areas with concomitant parallel or opposite changes of Chl and SST **B**) from Chl satellite and SST from the SODA reanalysis adapted from Martinez et al. (2009) and **D**) from Chl_{Svr-CCI} and SST_{NEMO}. The respective SST zero differences are shown on the maps as a thick black curve.

In their study, Martinez et al. (2009) analyzed two 5-year time periods apart each other by 15 years. They suggested that averaging observations separately over the two time-periods may have dampen the effect of interannual variability and reveal the decadal one. Most of the changes observed between the time periods covered by the two satellites are here confirmed based on the reconstructed Chlsvr-cci. However, the continuous 30-year time series of Chlsvr-cci provides new insights on the observed regime shifts (**Figure 12**). In the Pacific Ocean, the 1st EOF of Chlsvr-cci (**Figure 12A**) is close to the Chl spatial patterns obtained from the CZCS to SeaWiFS era (**Figure 11C**) and the PC remains highly correlated with the IPO over 1979-2010 (r=0.94 with p<0.001, **Figure 12B**). The Chl increase in the Indian Ocean, north-east of Madagascar toward the west

coast of Australia, between the 1980's and the 2000's also appears on the Chlsvr-CCI EOF. These temporal changes might also be related to the IPO variability (correlation between the IPO index and Chlsvr-CCI PC =0.6, p<0.001; **Figure 12C**).

In the Atlantic Ocean, CZCS-SeaWiFS Chl and Chlsvr-CCI 1st EOFs also share some similarities, including a decrease of Chl in the subtropical gyres and an increase in the equatorial/tropical regions. The associated PC (**Figure 12E**), exhibits a shift between 1979-1983 and 1998-2002 consistently with Figure 2C of Martinez et al. (2009). In this latter study, this change is attributed to a regime shift of the AMO. However, the AMO index is not correlated with the 1st Chlsvr-CCI PC (r=0.03, p=0.43) but rather with the 2nd mode (r = 0.43 with p=0.003), likely explaining the spatial discrepancies in **Figures 11A** *vs.* **11C**. Although the detailed analysis of Chl decadal variability is beyond the scope of the present study, these initial findings underscore the importance of continuous time series at regional/global scales to combine spatial and temporal information's and properly investigate Chl long-term variability.

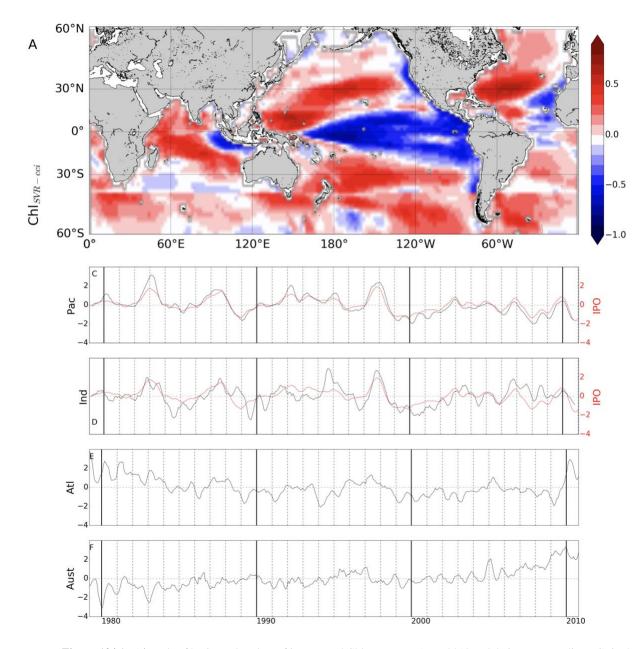


Figure 12 | **A**) 1st mode of basin-scale EOFs of interannual Chl_{Svr-CCI} over 1979-2010 and their corresponding PCs in the **B**) Pacific (23.2% of the total variance), **C**) Indian (15.2% of the total variance), **D**) Atlantic (13.5% of the total variance) and **E**) Austral Oceans (11.4% of the total variance). IPO is reported in red (right y-axis).

5 Summary and conclusions

In this paper, we assess the efficiency of a machine learning statistical approach based on support vector regression (SVR) to reconstruct surface Chl from oceanic and atmospheric variables. We first apply this strategy on self-consistent global dataset gathering physical predictors and Chl data simulated by a coupled physical-biogeochemical model simulation. Our results indicate that this nonlinear method accurately hindcasts interannual-to-decadal variations of the phytoplankton biomass simulated at global scale by the model, except at the boundaries of the subtropical gyres where the strong top-down control of zooplankton grazing in the numerical model is not accounted for by the SVR. Likewise, this statistical approach cannot yet reproduce Chl variability induced by nutrient inputs that are not directly related to our selected physical predictors, such as atmospheric iron deposit.

The SVR was then trained on satellite Chl observations. It accurately reproduces observed interannual Chl variations in most regions, including El Niño signature in the tropical Pacific and Indian Ocean as well as the main modes of Atlantic Chl variability. Despite an amplitude underestimation by half, it also accurately captures spatial patterns of Chl trends over the satellite period, with a Chl increase in most extratropical regions and a Chl decrease in the center of the subtropical gyres, as well as their changes between the CZCS and SeaWiFS era, while the model simulation generally fails to capture these long-term trends. Interestingly, while Chlpisces magnitude is closer to Chloc-cci than Chlsvr-cci, interannual variability and spatial trends of Chlpisces are farther than Chlsvr-cci to Chloc-cci. Equations representing the processes that govern the evolution of biogeochemical variables in a biogeochemical model are obviously less complex than the ones at play in the real ocean. We thus anticipated the modeled Chl to be easier to reconstruct than the satellite one. Additional complications were also expected through the

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reconstruction of satellite Chl from the model oceanic and atmospheric predictors, which may be less realistic than physical parameters derived from satellite measurements. As a consequence, the SVR is indeed slightly less efficient at reproducing the major satellite Chl patterns compared to the model ones but is surprisingly more efficient at capturing observed Chl temporal variations. This results in a normalized root mean square error generally weaker when reconstructing satellite data compared to the model one, although the predictors used are identical.

Machine learning techniques are powerful tools to statistically model non-linear processes. They require a significant amount of data to be trained and are well-suited to analyze remote sensing data. While several attempts have been made over the last decade to retrieve oceanic Chl content (Kwiatkowska and Fargion, 2003; Zhan et al., 2003; Camps-Valls et al., 2009; Jouini et al., 2013; Blix and Eltoft, 2018), the present work is one of the first attempt to use such machine learning techniques to reconstruct past time series of phytoplankton biomass at global scale. To our knowledge only Schollaert et al. (2017) tried to reconstruct the Chl multi-decadal variability in the tropical Pacific using a canonical correlation analysis built only from SST and SSH. Our SVR approach leads to higher correlations between reconstructed and satellite Chl in the tropical Pacific, highlighting the strength of such non-linear machine-learning methods with multiple predictors. These results emphasize deep learning approaches as promising tools to reconstruct multidecadal Chl time series in the global ocean, based on the knowledge of physical conditions. The successful use of surface variables only in reproducing Chl variability which is influenced by 3D-processes is here clearly noteworthy, and investigation of variable importance in the Chl reconstruction will deserve some future insights.

An obvious short-term perspective of the current study is to train a wider range of such statistical models with physical predictors derived from surface satellite observations but also

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observations within the water column which could be derived from Argo data (i.e., mixed layer and thermocline depth). Including complementary variables such as satellite particulate backscattering coefficient (as a proxy of the Particulate Organic Carbon) in the training/reconstruction process should also be considered. It would allow to investigate the extent to which the Chl variability reflects changes in phytoplankton biomass vs. cellular changes in response to light (e.g., Siegel et al., 2005; Westberry et al., 2008; Behrenfeld et al., 2015). The use of longitude and latitude as predictors may limit the ability to capture long-term trends in the evolution of the biogeochemical province boundaries, such as the expansion of the oligotrophic areas (Irwin and Oliver, 2009; Polovina et al., 2008; Staten et al. 2018). Thus, exploring deep learning schemes which may not explicitly depend on longitude and latitude, especially convolutional representations (Lecun et al., 2015), are particularly appealing. Further efforts need also to be dedicated to alleviate the issue of the underestimation of the long-term Chl trends. For instance, it would be noteworthy to investigate secular trends such as the 30% Chl decrease reported at global scale over the last century by Boyce et al. (2010), which remains largely debated (Mackas, 2011; McQuatters-Gollop et al., 2011; Rykaczewski and Dunne, 2011).

Whatever the methodology used (i.e. numerical models, satellite or *in situ* observations), they all have both advantages and drawbacks. *In situ* observations are considered as ground truth (with some errors/uncertainties depending for instance on the field measurement protocols), but are heterogeneous in time and space. Satellite Chl data provide a spatio-temporal synoptic view but they have their own measurements issues and uncertainties (e.g., sensor radiometric and spectral properties, atmospheric corrections, water constituents and their optical properties) and are limited to 20 years in their record length. Biogeochemical models are useful tools to (i) interpolate or extrapolate in time and space biogeochemical tracers such as Chl and to (ii)

investigate complex three-dimensional processes responsible for their variations. However, those models suffer from biases and are farther from *in situ* data than satellite observations. They are also not straightforward to run and require large computing resources. Thus, machine learning statistical schemes could be seen as a complementary tool to the "interpolate/extrapolate" use of biogeochemical models in providing a long-term synoptic surface view built from observations (being aware of the uncertainties associated with the variables used in the training schemes). Such methods, applied on observations only, will then provide an independent tool that may either question or enforce conclusions drawn from model simulations. Comparison between both methods and observations will also help to improve biogeochemical models with acute quantification of model biases and identification of the most meaningful predictors that may point to missing processes in biogeochemical models. As a conclusion, machine learning is a versatile tool that, associated with biogeochemical models and observations, may greatly enhance our view of global biogeochemistry.

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Authors contribution

EM led the project, analyzed results and wrote the 1st draft of the manuscript. TG provided physical model outputs. TG, ML provided support in the analysis and the writing of the manuscript. CF processed the machine learning approach with support from R. Sauzede. R. Fablet provided feedbacks on the statistical approach. All authors contributed to the development of the manuscript and provided feedbacks throughout its many stages of preparation.

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