Positive Indian Ocean Dipole events prevent anoxia off the west coast of India

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Abstract. The seasonal upwelling along the west coast of India (WCI) brings nutrient-rich, oxygen-poor subsurface waters to the continental shelf, favoring very low oxygen concentrations in the surface waters during late boreal summer and fall. This yearly-recurring coastal hypoxia is more severe during some years, leading to coastal anoxia that has strong impacts on the living resources. In the present study, we analyze a 1/4° resolution coupled physical–biogeochemical regional oceanic simulation over the 1960–2012 period to investigate the physical processes influencing the oxycline interannual variability off the WCI, that being a proxy for the variability on the shelf in our model. Our analysis indicates a tight relationship between the oxycline and thermocline variations in this region on both seasonal and interannual timescales, thereby revealing a strong physical control of the oxycline variability. As in observations, our model exhibits a shallow oxycline and thermocline during fall that combines with interannual variations to create a window of opportunity for coastal anoxic events. We further demonstrate that the boreal fall oxycline fluctuations off the WCI are strongly related to the Indian Ocean Dipole (IOD), with an asymmetric influence of its positive and negative phases. Positive IODs are associated with easterly wind anomalies near the southern tip of India. These winds force downwelling coastal Kelvin waves that propagate along the WCI and deepen the thermocline and oxycline there, thus preventing the occurrence of coastal anoxia. On the other hand, negative IODs are associated with WCI thermocline and oxycline anomalies of opposite sign but of smaller amplitude, so that the negative or neutral IOD phases are necessary but not the sufficient condition for coastal anoxia. As the IODs generally start developing in summer, these findings suggest some predictability to the occurrence of coastal anoxia off the WCI a couple of months ahead.

1 Introduction

The continental shelf off the west coast of India (WCI) is home to the largest coastal hypoxic system of the world ocean, spreading over an area of ∼180000 km² (Naqvi et al., 2000). These hypoxic conditions, characterized by an oxygen concentration lower than 20 µmol L⁻¹, occur in fall, right after the southwest monsoon. Importantly, substantial year-to-year changes in both the duration and intensity of this seasonal oxygen deficiency have been reported in the literature (e.g., Naqvi et al., 2009). While the oxygen concentrations in near-bottom waters are systematically low enough to trigger the conversion of oxidized nitrogen to molecular nitrogen, mostly through denitrification, this deficiency is even more severe in some years and the bottom waters turn sulfidic, a condition called coastal anoxia (Naqvi et al., 2006). These anoxic events have a tremendous impact on living resources (e.g., Diaz and Rosenberg, 2008), with more frequent episodes of fish mortality and a shorter fishing season, inducing a sharp decline in fish catches.

Observations from a series of ship cruises during September–October 1999 (Naqvi et al., 2000) off the WCI...
and time-series measurements from a fixed site off Goa since 1997 (Naqvi et al., 2009) indicate the occurrence of severe hypoxic conditions over almost the entire shelf and anoxic conditions close to the WCI, with the most intense anoxic event reported in fall 2001 and moderate ones in fall 1998 and 1999. In contrast, these data indicate that fall 1997 was characterized by far less hypoxic conditions. The frequent anoxic conditions which occurred during the 1998–2002 period were accompanied by a 3- to 5-fold decline in demersal fish catch in 1999 and 2001 compared to 1997. The total fish landing also remained low between 1998 and 2002, adversely affecting the economy based on fisheries, and the pelagic fish catches shifted from the dominance of mackerel to oil sardine from 1998 to 1999 (Krishnakumar and Bhat, 2008). There has been a remarkable revival of fisheries since 2003, apparently due to a relaxation of the intensity of oxygen deficiency, with no severe anoxic event reported over the recent years. Subsurface oxygen concentrations have also been reported to be significantly lower for the 1997–2004 period than for the 1971–1975 period (Naqvi et al., 2009). These observations thus reveal large interannual and decadal fluctuations in the oxygen deficiency along the WCI, but the processes responsible for these variations have not yet been understood (Naqvi et al., 2009).

As opposed to the other coastal hypoxic systems that have generally developed as a result of human activities (largely eutrophication) in the last few decades (Diaz and Rosenberg, 2008), the seasonal surface oxygen deficiency along the WCI is naturally driven. Indeed, it has been suggested that the oxygen-deficient conditions that develop in early fall along the WCI result from the seasonal upwelling, which brings poorly oxygenated subsurface waters from the oxygen minimum zone (OMZ) in the interior Arabian Sea (e.g., Morrison et al., 1999; Naqvi, 1987; Sarma, 2002) towards the surface along the continental shelf. This connection between the offshore and the shelf oxygen content has been discussed by several studies on the basis of in situ oxygen observations along different cross-shelf transects. For example, Banse (1959) showed a clear association between the seasonal upwelling and coastal hypoxia on the shelf off Cochin at 10° N in the last week of August 1957. A similar connection has also been reported by Carruthers et al. (1959) off Mumbai at 19° N in early November 1958 and later by Naqvi et al. (2009) off Mangalore at 13° N by late summer for several years. A recent study by Gupta et al. (2016) used 10 shelf transects near 10° N during 2012 to conclude that the upwelling of oxygen-deficient waters along the shelf break during the monsoon is the major process regulating the biogeochemistry on the shelf. Although these studies indicate a clear connection between the shelf oxygen content and upwelling at the shelf break, the actual processes in these exchanges are not well understood, owing to the lack of continuous and high-frequency observations.

The seasonal upwelling along the WCI starts in April, but the shallowest thermocline marking the peak of the upwelling is usually observed during September–October (e.g., Schott and McCreary, 2001). Local alongshore winds are, however, only favorable to upwelling during the southwest monsoon (i.e., they only have an equatorward alongshore component from June to August). This is an indication that upwelling along the WCI is to a large extent forced by remote winds (Shetye et al., 1990). Wind variations in the equatorial band and Bay of Bengal do indeed force coastal Kelvin waves that travel along the rim of the bay and up the WCI to influence the thermocline depth there (e.g., McCreary et al., 1993; Shetye, 1998). Recently, Suresh et al. (2016) demonstrated that wind variations in the vicinity of Sri Lanka are responsible for a large fraction of the seasonal upwelling along the WCI. By contrast with this seasonal variability of the upwelling and upper-ocean oxygen, there are currently little clues as to what causes the interannual variability along the WCI.

Identifying the main drivers of the near-surface oxygen interannual variations in regions of the main coastal hypoxic systems is an important endeavor as it may ultimately help to predict the occurrence of severe anoxic events. The large-scale climate modes have been suggested to influence the local oxygen variability in several coastal regions. For instance, the El Niño–Southern Oscillation (ENSO) strongly influences the oxygen concentrations along the coasts of Peru and Chile (Helly and Levine, 2004; Artz et al., 2006; Gutierrez et al., 2008), with intensified oxygenation associated with weak El Niño upwelling and intensified hypoxia associated with strong La Niña upwelling. As in the Pacific, the natural climate variability in the Indian Ocean could also be a potential candidate responsible for the near-surface oxygen interannual variations along the WCI, but that has so far not been evaluated. The main indigenous mode of Indian Ocean interannual climate variability is the Indian Ocean Dipole (hereafter IOD; Saji et al., 1999; Webster et al., 1999; Murtugudde et al., 2000). A “positive” IOD is characterized by a cooling and anomalously shallow thermocline in the eastern Indian Ocean and by a warming and anomalously deep thermocline in the central and western Indian Ocean, driven by anomalous easterlies in the central equatorial Indian Ocean. A “negative” IOD is associated with anomalous signals of opposite polarities. The IOD usually starts developing during boreal summer and peaks in fall (e.g., Saji et al., 1999). El Niño and La Niña events tend to induce, respectively, the positive and negative IODs in the Indian Ocean, but IOD can also occur independent of ENSO (e.g., Annamalai et al., 2003). The IODs induce larger amplitude variations in large-scale wind and thermocline depths than those associated with El Niño events over the Indian Ocean (e.g., Currie et al., 2013) and thus have the potential to affect the upwelling variations along the WCI through planetary wave propagation.

While the spatiotemporal density of observations in the eastern Pacific margin is adequate to accurately describe the monthly oxygen evolution along the west coast of South America over the past 3 decades (Helly and Levine, 2004;
Arntz et al., 2006), there is a dearth of long-term data from fixed sites in the Indian Ocean as compared to the Pacific and the Atlantic (Gupta et al., 2016), which in turn prevents building a reliable time series that could depict the year-to-year variations. On the other hand, three-dimensional coupled physical–biogeochemical models that include the oxygen cycle have contributed to improve the description and understanding of dynamics of hypoxic events in various coastal regions (Peña et al., 2010), such as the Gulf of Mexico (e.g., Hetland and DiMarco, 2008), Black Sea (e.g., Gregoire and Friedrich, 2004), and Baltic Sea (e.g., Eilola et al., 2009). Such models have also been used to investigate the mechanisms driving the spatial distribution (McCreary et al., 2013) and seasonal evolution of the OMZ in the interior Arabian Sea (Resplandy et al., 2012). There is, however, no dedicated modeling study to date, addressing the mechanisms that drive the interannual oxygen variability along the WCI. The present study aims at identifying the physical controls of the WCI interannual oxygen variability, with the help of a 40-year-long simulation from a 1/4° regional coupled physical–biogeochemical model. While the ∼25 km resolution of our model is insufficient to describe all the physical processes that influence the coastal upwelling (e.g., Huthnance, 1995; Allen et al., 2010) or the shelf–open-ocean interactions, we will show that our model still reproduces the essential characteristics of the large-scale oxygen fluctuations off the shelf break and that the offshore fluctuations provide a proxy for the variations on the shelf (Banse, 1959; Carruthers et al., 1959). Section 2 describes our model, data, and methods, and provides a brief evaluation of the model. Section 3 describes the main seasonal features of the thermocline and oxycline variability along the WCI in both the model and the observations. The strong influence of the IOD on the interannual oxycline variations along the WCI is then assessed from the model analysis in Sect. 4. Section 5 summarizes our results, discusses them in the context of earlier studies, and assesses the limitations of our approach.

2 Data and methods

2.1 Observations

We used the 1° resolution World Ocean Atlas 2013 (WOA13; Boyer et al., 2013) for evaluating the model accuracy at representing the large-scale climatological temperature and oxygen. The sea level is a good proxy for vertical movements of the thermocline in tropical regions (e.g., Fuku-mori et al., 1998), and hence we used the sea-level data from satellite altimetry produced by Ssalto/Duacs and distributed by AVISO (http://www.aviso.altimetry.fr/duacs/) as a proxy for the thermocline interannual variability along the WCI. In addition, we used the monthly level-3 Ocean Color Climate Change Initiative (OC-CCI) product, available at http://www.oceancolour.org/, that merges data from the Sea-WiFs, MERIS, and MODIS ocean color missions to evaluate the model surface chlorophyll climatological seasonal cycle. We also used oxygen measurements from the Candolim Time Series (CaTS) station located on the WCI on the inner shelf off Goa (∼15.5°N, 73.6°E) to construct a seasonal cycle of oxygen variations near the coast. This station has been established by the Council of Scientific and Industrial Research–National Institute of Oceanography (CSIR-NIO) and records the physical (temperature, salinity) and biogeochemical (oxygen, nitrate, nitrite, hydrogen sulfide, etc.) parameters from September 1997 onwards (see Maya et al., 2011, for a detailed description). This site lies approximately 10 km off the Candolim beach, where the depth of the water column is ∼28 m. Samples are taken at four depths: just below the surface, just above the bottom, and at two intermediate depths equally spaced between the surface and bottom. This dataset consists of a total of 142 vertical profiles during the 1997–2010 period and has been extensively used by previous studies (e.g., Naqvi et al., 2006, 2009, 2010a). Figure 1a shows the percentage of years sampled month-wise in this dataset. It indicates that in situ measurements have been performed at this station ∼50 % of the years over the 1997–2010 period for most calendar months, including September–October (the time of peaking of anoxic events), except for June (<10 %) and July (no data), when summer monsoon rough weather conditions prevent observations (Naqvi et al., 2006).
by five nutrients: NO$_3$, SiO$_4$, PO$_4$, SiO$_4$, and Fe. The ratios among C, N, and P are kept constant for the “living” compartments, at values proposed by Takahashi et al. (1985). The internal Fe contents of both phytoplankton groups and Si contents of diatoms are prognostically simulated as a function of ambient concentrations in nutrients and light level. Details on the red–green–blue model from which light penetration profiles are calculated are given in Lengaigne et al. (2007). The Chl / C ratio is modeled using a modified version of the photo-adaptation model by Geider et al. (1998). Dissolved oxygen is prognostic and evolves in response to physical conditions (advection, mixing), biological sources and sinks, and air–sea fluxes:

\[
\frac{\partial O_2}{\partial t} = \left( \frac{\partial O_2}{\partial t} \right)_{\text{Dyn}} + \left( \frac{\partial O_2}{\partial t} \right)_{\text{Bio}} + J_{\text{flux}},
\]

where \( \left( \frac{\partial O_2}{\partial t} \right)_{\text{Bio}} \) includes all biological processes affecting the concentration of O$_2$, \( \left( \frac{\partial O_2}{\partial t} \right)_{\text{Dyn}} \) accounts for large-scale and turbulent transport of oxygen and \( J_{\text{flux}} \) is the contribution of O$_2$ air–sea fluxes. The response of oxygen to biological processes \( \left( \frac{\partial O_2}{\partial t} \right)_{\text{Bio}} \) is computed as follows:

\[
\left( \frac{\partial O_2}{\partial t} \right)_{\text{Bio}} = \left( R_{1:0:c}^1 + R_{2:0:c}^2 \right) \left( \mu_{\text{NO}_3,P}^P + \mu_{\text{NO}_3,D}^D \right)
\]

\[
\text{New Production} + R_{1:0:c}^1 \left( \mu_{\text{NH}_4,P}^P + \mu_{\text{NH}_4,D}^D \right) - \lambda_{\text{DOC}} f(\text{O}_2) \text{DOC}
\]

\[
\text{Regenerated Production} - G^2 Z - G^M M - R_{0:0:c}^2 \text{Nitrification}
\]

\[
\text{Respiration} \quad \text{Nitrification}
\]

Oxygen is produced during net primary production (calculating the uptake of nitrate and ammonium by phytoplankton separately) by nanophytoplankton (P) and diatoms (D) and consumed by dissolved organic matter (DOC) remineralization, small (Z) and large (M) zooplankton respiration, and nitrification. This last term represents the conversion of ammonium into nitrate and is assumed to be photo-inhibited and reduced in suboxic waters. It is therefore a function of the ammonium and oxygen concentrations and photosynthetically available radiation. In this equation, the stoichiometric ratio \( R_{1:0:c}^1 \) represents the change in oxygen relative to carbon during ammonium conversion into organic matter, whereas \( R_{0:0:c}^2 \) denotes the consumption of oxygen during nitrification. Their values have been set, respectively, to 131 : 122 and 32 : 122 so that the typical Redfield ratio for oxygen is equal to 1.34 as proposed by Kortzinger et al. (2001).

When oxygen falls below a threshold value set to 6 umol L$^{-1}$, nitrate instead of oxygen starts to be increasingly consumed during the remineralization of organic matter, i.e., denitrification. To avoid negative oxygen concentration in the model, all processes consuming oxygen are switched off below a 10$^{-3}$ umol L$^{-1}$ threshold. Anammox is not represented in the model. At the bottom of the ocean, the model includes a very simple description of the sediment processes. The metamodel of Middelburg et al. (1996) is further used to compute the relative contribution of denitrification to the remineralization of the organic matter. Then, the flux of organic matter to the sediment is used to compute the oxygen and nitrate demands in the sediment, which are then imposed as boundary conditions to the model.

2.2 Model description

This study uses the NEMO (Nucleus for European Modelling of the Ocean; Madec, 2008) model that includes the PISCES (Pelagic Interaction Scheme for Carbon and Ecosystem Studies; Aumont et al., 2015) biogeochemical component. The PISCES model has 24 compartments, which include two sizes of sinking particles and four “living” biological pools, representing two phytoplankton (nano-phytoplankton and diatoms) and two zooplankton (microzooplankton and mesozooplankton) size classes. Phytoplankton growth is limited by five nutrients: NO$_3$, NH$_4$, PO$_4$, SiO$_4$, and Fe. The ratios among C, N, and P are kept constant for the “living” compartments, at values proposed by Takahashi et al. (1985). The internal Fe contents of both phytoplankton groups and Si contents of diatoms are prognostically simulated as a function of ambient concentrations in nutrients and light level. Details on the red–green–blue model from which light penetration profiles are calculated are given in Lengaigne et al. (2007). The Chl / C ratio is modeled using a modified version of the photo-adaptation model by Geider et al. (1998). Dissolved oxygen is prognostic and evolves in response to physical conditions (advection, mixing), biological sources and sinks, and air–sea fluxes:
The response of oxygen to dynamical processes is computed as

\[
\frac{\partial O_2}{\partial t}_{\text{Dyn}} = -u_H \cdot \nabla H O_2 - w \cdot \nabla v H O_2 + \nabla \cdot K_z \nabla Z O_2
\]

(3)

where \( u_H \) and \( w \) are, respectively, the horizontal and vertical currents and \( K_z \) is the vertical mixing coefficient computed by the dynamical model.

Finally, \( J_{\text{flux}} \) is the air–sea flux of \( O_2 (F_{O_2}) \) divided by the depth of the model surface layer, where

\[
F_{O_2} = k_w (\alpha O_{2\text{atm}} - O_2).
\]

(4)

\( k_w \) is the transfer velocity, \((\alpha O_{2\text{atm}} - O_2)\) is the difference in \( O_2 \) partial pressure between the air and surface seawater and \( \alpha \) is the solubility of \( O_2 \) in seawater.

The NEMO–PISCES coupled biophysical model has been successfully applied to various studies in the Indian Ocean (e.g., Koné et al., 2009; Resplandy et al., 2009; Currie et al., 2013; Keerthi et al., 2016), including the Arabian Sea OMZ (Resplandy et al., 2011, 2012). A more detailed model description is provided in the manuals for NEMO and PISCES, available online at http://www.nemo-ocean.eu/About-NEMO/Reference-manuals.

Here, we specifically use a 1/4° resolution (i.e., cell size \( \sim 25 \text{ km} \)) Indian Ocean regional configuration, which is a sub-domain of the global configuration described by Barnier et al. (2006). It has 46 vertical levels, with a resolution ranging from 5 m at the surface to 250 m at the bottom. The African continent forms the western boundary of the domain. The oceanic portions of the eastern, northern, and southern boundaries use radiative open boundaries (Treguier et al., 2001), constrained with a 150-day timescale relaxation to physical and biogeochemical inputs from a global simulation (Dussin et al., 2009). The circulation and thermodynamics of this regional configuration have been extensively evaluated and found to perform well in several Indian Ocean regions (Praveen Kumar et al., 2014), including the Arabian Sea (Nisha et al., 2013; Vialard et al., 2013; Keerthi et al., 2015, 2016) and the Bay of Bengal (Akhil et al., 2014, 2016).

The simulation starts from the rest and the WOA13 climatology temperature and salinity (Boyer et al., 2013). PISCES biogeochemical tracers are initialized from the WOA13 database for nutrients and a global simulation climatology for other tracers (Aumont and Bopp, 2006). After 5 years of spin-up with climatological surface and lateral boundary forcing, the model is forced with the Drakkar Forcing Set version no. 4.4 (DFS4.4; Brodeau et al., 2009) from 1958 to 2012. This forcing is a modified version of the CORE dataset (Large and Yeager, 2004), with atmospheric parameters derived from ERA40 reanalysis until 2002 (Uppala et al., 2005) and ECMWF analysis after 2002 for latent and sensible heat flux computation. Radiative fluxes are taken from the corrected International Satellite Cloud Climatology Project-Flux Dataset (ISCCP-FD) surface radiations (Zhang et al., 2004), while precipitation is specified from a blend of satellite products described in Large and Yeager (2004). All atmospheric fields are corrected to avoid temporal discontinuities and to remove known biases (see Brodeau et al., 2009, for details). In the following, the 1960–2012 period is analyzed.

2.3 Model climatology

The model’s ability to capture the climatological surface chlorophyll concentrations during the summer and winter monsoons is illustrated in Fig. 2. During the summer monsoon (Fig. 2a), seasonal blooms are observed along the coasts of the Arabian Peninsula and along the WCI in response to coastal upwelling that brings nutrients into the euphotic layer (e.g., Wiggert et al., 2005; Levy et al., 2007; Koné et al., 2009). The chlorophyll signal along the Somalia and Omani coasts extends offshore towards the central Arabian Sea (Fig. 2a) through offshore lateral advection of nutrients from upwelling regions, either by large-scale circulation or by eddy activity (e.g., Lee et al., 2000; Resplandy et al., 2011). During the winter monsoon (Fig. 2b), the cool, dry northeasterly winds in the Arabian Sea induce convective mixing and entrain nutrient-rich waters to the surface, triggering the chlorophyll bloom observed north of 15° N (Madhipratap et al., 1996). The model generally accurately reproduces these seasonal chlorophyll patterns (Fig. 2c and d).

As for observations, the largest chlorophyll bloom in summer occurs in the western Arabian Sea, with strong signals along the rim of the northern Indian Ocean, while winter is characterized by oligotrophic conditions in the southeastern Arabian Sea and higher concentrations in the northern and western parts. The largest discrepancy between modeled and satellite chlorophyll is an overall overestimation of the amplitude of the summer blooms, in the western Arabian Sea, east
of Sri Lanka and around the rim the Bay of Bengal (Fig. 2a and c). The seasonal chlorophyll patterns in fall are very similar although weaker than those in summer in both observations and the model (not shown).

The comparison of the modeled horizontal and vertical climatological oxygen distribution with that of WOA13 is shown in Fig. 3. In observations, the core of the OMZ is confined to the northern part of the basin (Fig. 3a) and expands vertically between 150 and 1000 m depth (Fig. 3b), with lowest subsurface oxygen concentrations found in the central and eastern part of the basin. The OMZ is thus shifted to the east of the region of highest biological production located along the west coast of the Arabian Sea (Fig. 2). The oxycline lies around ~100 m and is slightly shallower in the western and eastern part of the basin (Fig. 3b) because of the seasonal upwelling systems there. The model captures these observed oxygen patterns, with poorly oxygenated water confined to the northern Arabian Sea and highly oxygenated water found near the equator and farther south and an OMZ core located in the eastern part of the Arabian Sea despite a slight overestimation of the modeled oxygen content at depth in the Arabian Sea (Fig. 3b vs. d). We can also note that the model underestimates the depth of the OMZ core (~200 m in the model and ~300 m in observations). The upper-ocean vertical oxygen distribution is well captured, with a model oxycline depth of around 100 m (Fig. 3d), similar to the observed one (Fig. 3c).

2.4 Thermocline and oxycline depths

Both temperature and dissolved oxygen decrease with increasing depth below the mixed layer (Fig. 3b and d). The oxycline or thermocline depths are defined as the depths of maximum gradient. It is, however, customary to approximate those depths from a fixed isocontour, especially in tropical regions. Resplandy et al. (2012) used the depth of 100 µmol L$^{-1}$ as a proxy for the oxycline depth in their study, whereas Prakash et al. (2013) used 50 µmol L$^{-1}$. Here, we define the oxycline depth (hereafter OCD) as the depth of the 100 µmol L$^{-1}$ oxygen isocontour (following Resplandy et al., 2012) and the thermocline depth (hereafter TCD) as the depth of 23°C isotherm (following Prakash et al., 2013). The conclusions of our study are not sensitive to the above choices in the northern Indian Ocean (and especially in our focus region along the WCI), i.e., the results discussed below are similar when considering the 50 or 150 µmol L$^{-1}$ instead of 100 µmol L$^{-1}$, to define the oxycline depth and 20 or 25°C instead of 23°C to define the thermocline depth. The OCD and TCD were both derived from observed or modeled profiles using linear interpolation. To ascertain the variability along the WCI, we average relevant parameters over a box located between 10 and 15° N extending from the coast to 2° offshore (hereafter referred to as the WCI box, shown in Fig. 5). It must be noted that the results discussed below are not sensitive to any slight modifications of the box boundaries or offshore extent of this box. Interannual anomalies of all variables are calculated from monthly time series by subtracting the mean seasonal cycle and applying a 3-month smoothing to remove the sub-seasonal variations.

2.5 IOD index

To characterize the IOD variability, we used the standard definition of the Dipole Mode Index (DMI; Saji et al., 1999), which is calculated as the difference between the interannual sea surface temperature (SST) anomalies in the western (50–70° E, 10° N–10° S) and southeastern (90–110° E, 10° S–0°) equatorial Indian Ocean averaged over September–November. We have used the DMI derived from observed SST, but the very high correlation between the DMI based on model and observations (~0.94) makes our results fairly insensitive to the choice of either model or observationally based DMI. This index has been normalized by its standard deviation to make it dimensionless.

3 Results

3.1 Physical control of the seasonal oxygen variability off the WCI

The seasonality of the oxygen concentration within the core of the OMZ is very weak, but this is not the case for the OCD (Resplandy et al., 2012). Figure 4 shows the depth–time sec-
Figure 4. Seasonal evolution of oxygen (µmol L\(^{-1}\); color shaded) and temperature (°C; thin black contour) vertical profiles averaged over the WCI box (indicated as black frame in Fig. 5) from (a) WOA13 and (b) model. The oxycline and thermocline depths are marked by thick red and black lines, respectively, in both panels.

Figure 5. Maps of correlation between the mean seasonal cycle of oxycline and thermocline depths over the northern Indian Ocean from (a) WOA13 and (b) model. Values are masked when the oxycline could not be defined, i.e., when the oxygen concentration is above 100 µmol L\(^{-1}\) in the entire water column. The WCI box is marked as a black frame on each panel.

www.biogeosciences.net/14/1541/2017/ Biogeosciences, 14, 1541–1559, 2017

The strong dynamical control on the oxygen variability is further illustrated in Fig. 6, which displays spatial maps of observed and modeled OCD and TCD seasonal climatologies. During the spring inter monsoon (March–May), the TCD and OCD are spatially quite uniform and deep (∼100 m) in the southeastern Arabian Sea (Fig. 6a and e). This is also the case along the WCI south of 15°N despite the alongshore winds favorable to local upwelling, indicating a remote control of OCD and TCD variations there. In contrast, the shallower OCD and TCD near the southern tip of India (STI) during this season is consistent with upwelling favorable winds in this region (Fig. 6a and e). With the advent of the summer monsoon (June–August), the westerly monsoon winds drive a very strong offshore Ekman transport in the western Bay of Bengal and near the STI (Suresh et al., 2016), resulting in an upwelling signal, which shoals the OCD and TCD up to 60 m at the STI (Fig. 6b and f; Smitha et al., 2008; Gupta et al., 2016). Further north, the winds are almost perpendicular to the coast of western India (Suresh et al., 2016) and hence do not induce upwelling there. The shallow OCD and TCD signal from the STI propagates northward along the western Indian coast as an upwelling coastal Kelvin wave (Suresh et al., 2016; see Fig. 6b and f), with OCD and TCD patterns clearly suggestive of faster wave
Figure 6. Maps of the seasonal climatology of oxycline (m; shaded) and thermocline depths (m; contours with 10 m interval) from WOA13 during (a) March–May, (b) June–August, (c) September–November, and (d) December–February. (e–h) Same as (a–d) but from the model. Seasonal wind stress (N m$^-2$) patterns from the model forcing field are also shown as vectors in all panels.

The gaps in the CaTS observational dataset do not allow constructing a reliable time series of interannual upper-ocean oxygen content, to which our model could be validated. However, the data can still provide some estimate of the amplitude of interannual variability, which can be compared to that of our model (the whiskers in Fig. 7b and c indicate the amplitude of the variability around the mean seasonal cycle). Despite a slight underestimation in our model, the amplitude of interannual variability off the WCI is to a large extent driven by remote winds (Shetye et al., 1990). The wind near the STI is upwelling-favorable from about April to October and hence matches the seasonality of the upwelling of cold and low-oxygen waters, in agreement with the results of Suresh et al. (2016).
of the near-surface oxygen interannual variability is largest during SON, both in the shelf observations and the model. This further corroborates the offshore–shelf connection discussed above.

A seasonally shallow OCD combined with a larger interannual variability in fall creates a window of opportunity for the occurrence of coastal anoxic events. Figures 7d and e display the monthly percentages of the occurrence of hypoxic profiles from CaTS (on the shelf) data and the model (offshore). While the general patterns of oxygen (oxycline) variability on the shelf and offshore of the coast remain similar, the actual upper-ocean oxygen content and vertical oxygen profiles are different. Hence, we have used different thresholds to detect hypoxic profiles in the observation and the model. Consistent with previous literature, anoxic events are most likely to occur from August to November in the model and the shelf data, as expected from the very shallow background oxycline at that time of the year. This justifies our focus on the fall period for analyzing the processes that drive the modeled interannual variability of the WCI oxycline in the following.

We previously demonstrated a tight relationship between the seasonal variability of OCD and TCD in the eastern Arabian Sea. Figure 8a exhibits a similar relation in large portions of the northern Indian Ocean for fall interannual OCD and TCD anomalies in the model. A comparison with observations is unfortunately not feasible due to lack of a basin-scale dataset for interannual OCD anomalies. The cor-
The influence of remote forcing at WCI is further established in Fig. 9. Due to the unavailability of the continuous oxygen observations (see Sect. 2.1), we cannot directly evaluate the modeled oxygen interannual variability in the WCI region. However, we evaluate the modeled TCD interannual variability, which is closely tied to the OCD interannual variability (∼0.95 correlation, Fig. 8b). The modeled interannual TCD anomalies in the WCI box (red curve in Fig. 9a) agree well (0.84 correlation) with sea-level interannual anomalies (a good proxy for TCD variations in stratified regions) from altimeter measurements (blue curve in Fig. 9a) during fall. Despite instances, when the model agreement is weaker, like during the 2002–2006 period, both the modeled TCD and the observed sea level indicate the strongest thermocline shoaling in fall 1999 and a deeper than usual thermocline in fall 1994, 1997, and 2008. We will exploit this ability of our model to capture the observed TCD interannual variations along the WCI to further examine the processes responsible for the OCD interannual variability. Along with fall TCD and sea-level anomalies at WCI, Fig. 9a also displays the fall interannual anomalies of zonal winds at the STI (black dashed...
The relationships between the modeled interannual variability of the OCD along the WCI and that of the OCD, TCD, SST, and wind on the basin scale are demonstrated in Fig. 10a and b, which show regression maps of fall interannual anomalies of these variables to the time series of the WCI OCD anomalies shown in Fig. 8b (black line) over the 1960–2012 period. These maps display the typical basin-scale anomalies corresponding to an anomalously deep OCD in the WCI. Consistent with Fig. 9, the fall WCI OCD variations are not merely local but are associated with basin-scale ocean–atmosphere interannual anomalies over the entire equatorial and northern Indian Ocean. An anomalously deep OCD off the WCI is usually associated with deeper OCD and TCD (i.e., positive anomalies) in the southeastern Arabian Sea and in the vicinity of Sri Lanka and the STI (Fig. 10a and b). Positive OCD anomalies off the WCI are also related to shallower OCD and TCD in the eastern Indian Ocean and along the eastern rim of the Bay of Bengal (i.e., negative anomalies). The associated large-scale wind patterns (Fig. 10a) explain these interannual OCD and TCD patterns. Anomalous easterlies in the equatorial band force upwelling equatorial Kelvin waves that shoal the OCD and TCD in the EEIO. These signals further propagate around the rim of the bay as upwelling coastal Kelvin waves, thereby shoaling the TCD and OCD there. The strong negative correlation between the WCI OCD and STI zonal winds interannual fluctuations (−0.73 over the entire period; see Fig. 11b) further illustrates the strong influence of these winds in driving OCD and TCD interannual fluctuations in the southeastern part of the Arabian Sea. In contrast, the local alongshore wind variations along the WCI have a weaker influence on the WCI OCD, with a negative correlation of −0.25 (Fig. 11c), confirming the dominance of remote forcing from the STI region over the influence of local winds.

3.3 The specific role of the IOD

The SST variations associated with the OCD signals in the WCI region are characterized by a clear signal in the EEIO (near the Sumatra coast) and weaker signals of opposite sign in the western Indian Ocean. As pointed out before, the patterns shown in Figs. 9b, c and 10a, b are reminiscent of the IOD signature (Saji et al., 1999; Webster et al., 1999; Murtugudde et al., 2000), an Indian Ocean coupled ocean–atmosphere climate mode that peaks in fall, as discussed in the introduction. This is further demonstrated in Fig. 10c and d, which display regression maps of interannual anomalies of OCD, TCD, SST, and winds onto the boreal fall DMI. The resulting patterns, representing the typical anomalies associated with a positive IOD phase, are strikingly similar to those displayed in Fig. 10a and b (pattern correlation of ~0.85). This highlights the strong link between the IOD events and the WCI oxycline year-to-year variations. We further examine the relationship between the WCI OCD and the IOD in Fig. 11a, which displays time series of fall OCD interannual anomalies and the fall DMI (computed from both modeled and observed SST, the correlation between them being 0.94). Consistent with the regression map, we find a high correlation between the WCI OCD and the

![Figure 10. Regression patterns of fall interannual anomalies of modeled (a) oxycline (m; shaded) and wind stress (N m$^{-2}$; vectors) and (b) thermocline (m; shaded) and SST (${}^\circ$C; contours with 0.1 $\circ$C interval) onto the fall oxycline interannual anomalies averaged over WCI box normalized by its standard deviation. (c–d) Same as (a–b) but regressed onto the observed fall DMI index.](image-url)
DMI (~0.67/0.62 with modeled and observed DMI, respectively).

The influences of positive and negative phases of IOD on the WCI OCD (and TCD, not shown) are, however, not symmetrical (Fig. 11a): most of the positive IOD events cause a deepening of OCD along WCI (e.g., 1961, 1967, 1994, 1997), while negative IOD can either be associated with a shoaling (e.g., 1996, 1998, 2010) or even a deepening (e.g., 1979–1981). To further illustrate this asymmetry, Fig. 12 provides a scatterplot of the fall DMI versus the fall interannual anomalies of OCD off the WCI. This scatterplot confirms that there is an asymmetric impact of positive and negative IODs on the OCD along the WCI. The WCI OCD response to positive IODs is very robust, with all positive IOD events (DMI > 1) except one being systematically associated with a deepening. The WCI response to negative IOD is weaker and much less systematic. As discussed above, negative IOD events (DMI < −1) are generally related to negative OCD anomalies but can also be related to positive OCD anomalies (Fig. 11a). Figure 13 provides a composite of the temporal evolution of the anomalous OCD and TCD along the WCI along with zonal wind stress variations at the STI for signal associated with positive/negative IOD events. It must be noted here that time series for negative IOD events has been multiplied by −1 to ease comparison with positive events. Figure 13a and b illustrate again the 2 to 3 times weaker response of the WCI OCD for negative compared to positive IOD events. This weaker response is related to slightly weaker zonal wind anomalies at the STI (Fig. 13c) that consequently trigger a weaker coastal Kelvin wave response along WCI and thus weaker WCI TCD and OCD anomalies. It is therefore likely that the weaker and less robust wind signal associated with negative IODs at the STI compared to positive IODs may explain part of the asymmetry seen in Fig. 12. We will discuss the possible causes for this asymmetry in Sect. 4.2.

4 Summary and discussion

4.1 Summary

The year-to-year variations of coastal hypoxia along the west coast of India (WCI) have been identified by Naqvi et al. (2000, 2009), along with their strong impacts on fisheries and the ecosystem. The mechanisms controlling these variations have, however, not yet been elucidated. The present study offers new insights on the physical controls of coastal hypoxia along the WCI. To that end, we used an
V. Parvathi et al.: IOD influence on anoxia off the WCI

Figure 13. Seasonal evolution of anomalous composites of (a) WCI OCD, (b) WCI TCD, and (c) STI zonal wind stress during positive (red) and negative (blue) IOD events. Positive (negative) IODs are defined when the DMI averaged for the fall season is greater (lesser) than 1 °C. The whiskers indicate the 95 % confidence interval on the composited value. Positive IOD events considered in this composite are the years 1961, 1963, 1967, 1972, 1977, 1982, 1994, 1997, and 2006, while negative ones are 1964, 1973, 1974, 1975, 1979, 1981, 1984, 1992, 1996, 1998, and 2010. Negative IOD composite time series has been multiplied by −1 to ease comparison with positive events.

The modeled and observed offshore climatological seasonal cycles of oxygen match in situ measurements on the shelf, with a strongest seasonal oxygen deficiency and highest occurrence rate of anoxic events during boreal fall. It is suggested that the upwelling of oxygen-depleted subsurface waters at the shelf break influences the occurrence of anoxic events over the western Indian continental shelf.

The shallow oxycline in fall combines with a large interannual variability at this time of year to create a window of opportunity for coastal anoxic events. Our model analysis further indicates that there is a tight coupling between the thermocline and oxycline variability in this region on both seasonal and interannual timescales, indicative of a strong physical control of the oxygen variability through vertical advection. Interannual thermocline fluctuations along the WCI are related to basin-scale wind, thermocline, and oxycline depth perturbations associated with IOD events, an Indian Ocean coupled ocean–atmosphere climate mode that peaks in fall. Positive IOD events are associated with easterly wind anomalies in the central equatorial Indian Ocean and extend meridionally up to the southern tip of India. These easterly wind anomalies trigger downwelling coastal Kelvin waves that propagate along the WCI and deepen the thermocline and oxycline in boreal fall, thereby preventing the occurrence of coastal anoxia off the WCI during positive IOD events. Our model results also suggest an asymmetry between the impact of positive and negative IOD events on the WCI oxycline depth. The westerly wind anomalies at the southern tip of India do indeed have a smaller amplitude during negative IODs than their easterly counterparts during positive IODs, thus resulting in a weaker and less consistent shoaling of oxycline and thermocline along WCI during negative IOD events.

4.2 Discussion

4.2.1 Influence of the IOD on the interannual oxygen variability along the WCI

Previous studies have demonstrated the impact of large-scale climate modes on year-to-year variations of the oxygen deficiencies in coastal hypoxic systems elsewhere in the world ocean. In the Pacific, El Niño conditions lead to intensified oxygenation along the coasts of Peru and Chile as a result of weak upwelling (e.g., Arntz et al., 2006; Gutierrez et al., 2008), while in the Atlantic, the Benguela Niño leads to intensified anoxia along the Namibian shelf (Monteiro et al., 2008). The western continental shelf of India is home to the largest naturally formed coastal hypoxic system in the world. In this study, we identify, for the first time, the IOD as the major climatic driver of the year-to-year oxycline and thermocline variations offshore of the WCI. Though the IOD has a weaker thermocline depth signature on the west than on the east coast of India, it has stronger societal consequences as it influences the WCI seasonal upwelling that brings suboxic
winters very close to the surface during fall. Although the IOD influence on the west Indian coast has never been reported so far, it has regularly been reported in the in the Bay of Bengal. In line with our results, Aparna et al. (2012) did indeed show that IOD events drive strong sea-level and thermocline fluctuations along the rim of the bay in fall, through coastal Kelvin wave propagation from the equatorial region. Akhil et al. (2016) further demonstrated that this remote forcing also drives counterclockwise anomalous horizontal currents in fall in the bay, which in turn leads to large interannual variations of sea surface salinity in the southern Andaman Sea. On the biogeochemical side, Wiggert et al. (2002, 2009) and Currie et al. (2013) demonstrated that IOD events are responsible for large interannual chlorophyll variations in the southeastern Bay of Bengal and at the STI. Finally, the IOD signature found in the Arabian Sea in the present study has already been described for the Bay of Bengal in terms of sea level (Aparna et al., 2012) and chlorophyll (Currie et al., 2013).

The influence of IOD is further shown to be larger for its positive than its negative phase. Our results suggest that part of the weaker WCI oxycline depth response during negative IOD may be explained by the weaker wind stress anomalies at the STI associated with negative IOD events. This weaker wind amplitude could simply be related to the tendency of negative IOD events to be weaker than their positive counterpart (Saji and Yamagata, 2003; Hong et al., 2008; Cai et al., 2013) or to asymmetries in the spatial patterns of winds associated with the nonlinear response of deep atmospheric convection to SST anomalies of each phase of the IOD. A more precise understanding of this asymmetry would require an in-depth investigation of the processes that control the wind variations at the STI and the thermocline along the WCI in response to positive and negative IOD events.

Our findings partly explain the substantial year-to-year changes in both the duration and intensity of the observed seasonal oxygen deficiency over the western Indian shelf (Naqvi et al., 2009). None of the anoxic events reported by Naqvi et al. (2009) (black stars in Fig. 12) lies in the upper right quadrant of the scatterplot shown in Fig. 12, indicating that positive IODs systematically prevent the occurrence of anoxic events. For instance, the relaxation of anoxic conditions in early fall 1997 reported by Naqvi et al. (2009) is in line with the occurrence of very strong positive IOD during that year. Most anoxic events are found in the lower left quadrant, i.e., near neutral or negative IOD conditions and an anomalously shallow offshore oxycline. Neutral or negative IOD years are, however, not necessarily anoxic, indicating that a neutral or negative IOD is a necessary but not a sufficient condition for severe anoxia. A recent study by Gupta et al. (2016) revealed that the oxygen deficiency in 1959 along the WCI was more severe than in 2012, a conclusion consistent with the occurrence of a negative IOD in 1959 and a positive one in 2012. Similarly, in situ measurements also revealed that subsurface oxygen concentrations were significantly lower at the turn of the 20th century than in the 70s (Naqvi et al., 2009): our simulation exhibits a similar behavior (see Fig. 11a), showing many years with shallower than normal OCD in the later period and systematically deeper than normal OCD during 1970s. The causes for those decadal variations need to be investigated in greater detail.

The ∼ 0.7 correlation between IOD variability and oxycline variations along the WCI implies that ∼ 50% of the interannual oxycline variance is explained by the IOD at this location. This relationship between the IOD and year-to-year variations of seasonal anoxic conditions along the shelf may facilitate an advance warning for the possible occurrence of severe anoxic events. Recent studies do indeed indicate that skillful predictions of mature IOD events in fall can be achieved about one season ahead (e.g., Luo et al., 2007; Wang and Justic, 2009; Zhao and Hendon, 2009; Sooraj et al., 2012) and up to two seasons ahead in the case of large IOD events (Luo et al., 2007, 2008; Shi et al., 2012). Those predictions of IOD events should allow providing a warning about the likelihood of severe anoxic conditions along the WCI during spring or summer. A predicted positive IOD is indeed associated with very low chances of such an anoxic event, while neutral or negative IOD conditions may be associated with the occurrence of such an event.

It must, however, be kept in mind that other factors are also likely to contribute to the reported interannual fluctuations of hypoxic conditions in this region. Naqvi et al. (2009) for instance suggested that increased productivity due to increased nutrient loading from land associated with anthropogenic activities might have the potential to trigger a shift from natural suboxic to anthropogenic anoxic conditions during recent decades. This hypothesis, however, cannot explain the relaxation of the intensity of oxygen deficiency in recent decades. Another contributing factor could be related to changes in local hydrographic variations. For instance, interannual variations of the land runoff along the Western Ghats, local precipitation during the summer monsoon, or input of Bay of Bengal freshwater during the northeast monsoon (e.g., Jensen, 2001) could modulate the upper-ocean haline stratification, ventilation of the subsurface waters, and hence the subsurface oxygen content along the WCI. Finally, local along-shore wind variations may modulate the intensity of coastal upwelling and hence the amount of oxygen-depleted waters brought to the shelf. The influence of these factors thus requires further investigation.

4.2.2 Model limitations

An obvious limitation of the current study is the spatial resolution of our model (∼ 25 km). While our model has a reasonable representation of the temperature and oxygen seasonal variations in the deep ocean off the shelf break, its spatial resolution is not sufficient to resolve the details of physical processes controlling the upwelling along the shelf break (e.g., Huthnance, 1995; Allen et al., 2010). For the case of
the narrow continental shelf along the west coast of North America, several studies have shown that a minimum horizontal resolution of 10 km is required (Marchesiello et al., 2003; Veneziani et al., 2009). For example, at 1/4° resolution our model is eddy-permitting but not eddy-resolving, and hence does not fully capture oceanic mesoscale eddies, which play an important role in the exchanges between the shelf and the open ocean in upwelling regions (e.g., Marchesiello et al., 2003; Bettencourt et al., 2015; Vergara et al., 2016). Another limitation arises from the absence of tidal forcing in the model, which may play an important role, as strong internal tides can be generated on the shelf break and contribute to enhancing the thermocline vertical excursion and mixing, which can both contribute to bringing more deep-ocean oxygen-deficient water to the shelf (Monteiro et al., 2008).

While our model does not reproduce the details of exchanges between the shelf and open ocean, we have just used it as a proxy of the behavior of open ocean, off the WCI. Several studies have already pointed towards the influence of offshore oxygen variations on the variability of hypoxic conditions along other coastal regions (e.g., Grantham et al., 2004; Helly and Levine, 2004; Arntz et al., 2006; Gutierrez et al., 2008). As was shown in Figs. 6 and 7b, the model and WOA climatology vertical oxygen distributions agree quite well, both in terms of the oxycline depth and near-surface value. The CaTS data, on the other hand, is representative of what happens much closer to the coast and displays much lower oxygen levels than seen further offshore in WOA and the model. This may of course partially be due to shortcomings in representing physical exchanges between the shelf and open ocean at the current resolution of our model and existing oxygen dataset in the region. But biological processes are also known to be a prominent oxygen consumption factor on the shelf, in particular in the benthic zone where the enhanced concentration of particulate matter above sediments is associated with a high oxygen demand (e.g., Cowie, 2005).

The crude parameterization of sediments in the model probably does not consume enough oxygen very close to the coast. On the other hand, the good phasing between the oxygen seasonal variability offshore (in the model and WOA) and shelf (CaTS) data (Fig. 7b and c) suggests that the offshore variability is probably an important driver of the oxygen content on the shelf. However, a proper representation of benthic biological processes would probably be needed to represent the low oxygen levels very close to the coast (Fig. 7c). Dedicated studies at a higher spatial resolution with sensitivity tests on the representation of nearshore biological processes will probably be needed in order to better understand how the representation of nearshore biological processes constrains the coastal oxygen representation.

4.2.3 Observational requirements

On the observational front, the current spatiotemporal sampling does not allow building reliable long-term time series of the month-to-month oxygen variations along the shelf and offshore. Despite the establishment of frequent measurements of the oxygen profile off Goa since September 1997, the numerous unsampled months (July and August are almost unsampled because of rough weather conditions) and the strong sub-monthly variability prevent a continuous monitoring of oxygen variations along the WCI. A reasonable number of moorings or Argo drifters with oxygen and temperature sensors along the shelf and further offshore would allow a finer description of the oxygen variability and of its relationship with temperature and connection with the offshore variations. In order to establish unequivocal evidence for the shelf–open-ocean interactions, future studies should also consider improved observations such as repeated glider transects or triad of moorings (shelf, shelf break, and open ocean) monitoring both physical and biogeochemical quantities in this region.

4.2.4 Hypoxia in other regions of the northern Indian Ocean

Though the present study is focused on the WCI, our Indian Ocean configuration model allows assessing other regions where near-surface hypoxia can occur. Figure 14 shows the percentage of profiles where oxygen concentrations below 80 µmol L$^{-1}$ occur within the top 50 m. This threshold is indicative of the limit under which many organisms start to suffer from physiological stress that could ultimately lead to death (Vaquer-Sunyer and Duarte, 2008). This analysis indicates that the coast of Oman can also experience hypoxic...
conditions as reported in the literature (e.g., Piontkovski and AL-Oufi, 2015), although hypoxia along Oman is never as severe as that off WCI (Naqvi et al., 2010b). Figure 14 indicates that the northwestern Bay of Bengal can also experience near-surface low-level oxygen concentrations, as reported from a series of ship cruise measurements by Sarma et al. (2013). Further examination of the mechanisms driving these hypoxic events reveals that the IOD strongly impacts the oxygen variability in the northwestern Bay of Bengal: positive IOD events generally inducing a shoaling of the oxycline in this region (see Fig. 10c) through upwelling coastal Kelvin wave propagation from the equatorial region. In contrast, the influence of IOD along the Omani coast is almost negligible (Fig. 10c), and oxygen variations here seem to be related to offshore Ekman pumping (not shown). Further dedicated studies are needed to better understand the oxygen variability in these sensitive regions and their potential impacts on the ecosystem and fisheries.

Data availability. The (CaTS) data archived at the CSIR-NIO data center and Chemical Oceanography Division have been used in Figs. 1 and 7. These datasets are not publicly available. Hence we do not provide the data or any access link. All other datasets are publicly available: WOA13 – https://www.nodc.noaa.gov/OC5/woa13/woa13data.html (NOAA, 2015); SLA – http://www.aviso.altimetry.fr/en/data/products/sea-surface-height-products/global.html (AVISO, 2014); OC-CCI – https://www.oceancolour.org/ (Plymouth Marine Laboratory – PML, 2014); NEMO model source codes are downloadable after registering at http://www.nemo-ocean.eu/About-NEMO.

Competing interests. The authors declare that they have no conflict of interest.

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