Seasonal ITCZ migration dynamically controls the location of the (sub)tropical Atlantic biogeochemical divide

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Inorganic nitrogen depletion restricts productivity in much of the low-latitude oceans, generating a selective advantage for diazo-
trophic organisms capable of fixing atmospheric dinitrogen (N2). However, the abundance and activity of diazotrophs can in turn be controlled by the availability of other potentially limiting nutrients, including phosphorus (P) and iron (Fe). Here we present high-resolution data (0.3°) for dissolved iron, aluminum, and inorganic phosphorus that confirm the existence of a sharp north-south biogeochemical boundary in the surface nutrient concentrations of the (sub)tropical Atlantic Ocean. Combining satellite-based precipitation data with results from a previous study, we here demonstrate that wet deposition in the region of the intertropical convergence zone acts as the major dissolved iron source to surface waters. Moreover, corresponding observations of N2 fixation and the distribution of diazotrophic Trichodesmium spp. indicate that movement in the region of elevated dissolved iron as a result of the seasonal migration of the intertropical convergence zone drives a shift in the latitudinal distribution of diazotrophy and corresponding dissolved inorganic phosphorus depletion. These conclusions are consistent with the results of an idealized numerical model of the system. The boundary between the distinct biogeochemical systems of the (sub)tropical Atlantic thus appears to be defined by the diazotrophic response to spatial–temporal variability in external Fe inputs. Consequently, in addition to demonstrating a unique seasonal cycle forced by atmospheric nutrient inputs, we suggest that the underlying biogeochemical mechanisms would likely characterize the response of oligotrophic systems to altered environmental forcing over longer timescales.

Significance

Low concentrations of fixed nitrogen restrict phytoplankton growth in much of the low-latitude surface oceans. Diazotrophic cyanobacteria are capable of fixing atmospheric dinitrogen, thereby replenishing the overall pool of fixed nitrogen. As a consequence, spatial–temporal variability in diazotrophy can potentially influence the global nitrogen cycle. Here we show that movement in the region of elevated iron concentrations tied to the seasonal migration of the intertropical convergence zone drives a shift in the latitudinal distribution of dinitrogen fixation and corresponding phosphate depletion in surface waters. Surface nutrient concentrations and diazotrophic activity divide the (sub)tropical Atlantic into a high-phosphate, low-iron system in the south, and a low-phosphate, high-iron system in the north.


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The boundary region between these (sub)tropical gyre systems is typically characterized by high rates of N₂ fixation, which are dominated by the diazotrophic cyanobacterium *Trichodesmium* spp. (12, 29–31), and appear to be spatially correlated with some of the areas of highest DFe concentrations (12). Surface nutrient concentrations and diazotrophic activity thus appear to define a biogeochemical division of the Atlantic between a high DIP, low DFe (HPLFe) system in the south, and a low DIP, high DFe (LPHFe) system in the north (12).

Although the biogeochemical division of the (sub)tropical Atlantic has previously been interpreted on the basis of the hypothesized Fe control of diazotrophy (12, 13), there remains the possibility that the observed spatial patterns could have other forcing factors (32). Moreover, the wider regional and global importance of Fe controls on diazotrophy in both the modern and paleo ocean remains a matter of active debate (12, 17, 33–35) with important implications for our understanding of, for example, glacial/interglacial nitrogen and carbon cycling (1, 36–38). More direct tests of the Fe–diazotrophy limitation hypothesis are thus desirable. In particular, predictable large-scale dynamic responses of marine diazotrophy and subsequent P cycling to seasonal variability in external Fe inputs has not previously been described.

**Results and Discussion**

The Sahara and Sahel deserts deliver large quantities of mineral dust to the (sub)tropical North Atlantic (22, 39). Tropical Rainfall Measuring Mission (TRMM) and Moderate Resolution Imaging Spectroradiometer (MODIS) satellite data for the period of our research cruise (February–March 2011), indicated that the northern part of the study region received significant dry (Fig. 1A) and wet (Fig. 1B) atmospheric deposition fluxes. However, the tropical South Atlantic was strongly shielded from desert dust by the InterTropical Convergence Zone (ITCZ), which had its core at ∼1°N during our cruise period, close to the southernmost latitude reached during its annual migration (40, 41) (Fig. 1B).

In contrast, during a previous study in the region (October–November 2005; Atlantic Meridional Transect [AMT-17]; ref. 42) the ITCZ was located at ∼6°N, nearer to the northernmost extent (Fig. 1C).

The low pressure system of the ITCZ, formed by rapidly rising humid air (41), is characterized by heavy precipitation of up to ∼80 mm h⁻¹ (45) (Fig. 1B and C). Aerosols are rapidly scavenged by this band of precipitation (24, 44), and the ITCZ system also restricts southward transport of desert-dust-rich air masses, as was evidenced by airmass back trajectories (HYbrid Single-Particle Lagrangian Integrated Trajectory [HYSPLIT]) calculated for the periods of both cruises (Fig. S1). Precipitation has a major influence on sea surface salinity (SSS) in the tropical Atlantic, with the latitudinal migration of the ITCZ (40, 41) resulting in a strong seasonal cycle in local salt storage in the region between ∼0° and 10°N (45, 46). Correspondingly, we observed that the core of the ITCZ was associated with SSS minima during both meridional transects in 2005 and 2011 (Fig. 2).

Wet deposition within the ITCZ has previously been proposed to dominate trace metal inputs in the region (24, 47). Correspondingly, we observed that dissolved aluminum (DAI), a non-nutrient tracer of atmospheric inputs (24, 48) and the micronutrient DFe were highly correlated both with each other and with SSS along our two high-resolution 2011 transects (Fig. 2). The relationship of DFe and DAI with SSS for the earlier 2005 transect was remarkably similar (Fig. S2), despite a ∼4° latitudinal offset in the region of minimum SSS and maximum DFe and DAI concentrations.

Tighter correlation between DFe and precipitation (Fig. 1B) and SSS (Fig. 2), rather than aerosol optical thickness (Fig. 1A), suggest that wet deposition dominated Fe input in this region. More efficient solubilization of Fe from mineral dust particles during wet deposition (49, 50) due to a low rainwater pH (pH 4–6), the presence of Fe-binding ligands (51), and continuous photochemical reactions producing soluble Fe(II) in rain droplets (49) may all contribute to such dominance. Our data (Fig. 2), together with previous observations (12, 24, 47), thus strongly argue for wet deposition associated with the ITCZ forming the major source of DFe to the surface waters of the tropical Atlantic, as the ITCZ simultaneously shields the South...
Atlantic from atmospheric dust inputs, resulting in low DFe concentrations (12, 28).

Sharp gradients in surface water DIP corresponded with the southern boundaries of the regions of minimum SSS and enhanced DFe in 2011 (Fig. 2). The inverse relationship between DFe and DIP thus agreed with our 2005 observations (12); however, the strong gradient region was offset to the south by 5°–10°, corresponding to the seasonal southward displacement of the ITCZ (Fig. 3). Moreover, observed N₂ fixation rates, alongside the abundance of *Trichodesmium*, peaked in the broad transition region between the HPLFe waters to the south and the LPHFe waters to the north in 2011 and 2005, thus demonstrating an apparent seasonal latitudinal shift consistent with previous data from the region (52). Consequently, data from three crossings of the ITCZ (Figs. 2 and 3) were fully consistent with our proposed conceptual model, whereby spatiotemporal variability in Fe inputs can drive variability in diazotrophy and subsequently control the availability of excess DIP in a system (12, 13).

To further illustrate the plausibility of the proposed mechanism, we expanded a widely used simple numerical model (55–55) to incorporate an idealized two-layer (surface–thermocline) horizontal transport framework, which can be taken to represent interactions between differential nutrient supplies (N, P, and Fe) and (non)diazotrophs along any generic upper-ocean circulation pathway (SI Materials and Methods). In the current case, our model configuration can be considered an idealized representation of the upper limb of the Atlantic overturning circulation (12, 56), where surface and thermocline waters apparently accumulate geochemical and isotopic imprints of N₂ fixation during this large-scale northward transport (12, 57). Although simple, the underlying functional equations (55) are effectively identical to those used within more complex biogeochemical/ecosystem general circulation models (58, 59). In consequence, the qualitative behavior is expected to be consistent with more complex circulation and ecosystem scenarios, and the responses to variable forcing will be more fully tractable.

Qualitative steady-state outputs from our model setup were highly insensitive to parameter choices within well-defined bounds (Table S1). Thus, running the model with an initial subsurface excess of DIP, as occurs almost ubiquitously throughout the global (sub)tropical oceans (4, 55), and a spatially varying Fe input, specifically a peak (or equivalently a jump) in input, resulted in different areas of the model domain corresponding to two well-defined regimes. Specifically, a “switch” occurs between a modeled initial HPLFe and subsequent LPHFe condition (Fig. S3), with the boundary region of this switch, a biogeochemical divide, being related to the external Fe input and associated with a corresponding peak in diazotrophy (Fig. S4). Formally, this occurs because diazotrophy increases sharply at the location in the model domain where the ratio of total Fe:DIN inputs is sufficient to maintain surface DFe concentrations above the corresponding equilibrium resource concentration for diazotrophs, typically denoted *R*ₘ (55).

As previously noted (55), steady-state model outputs were highly analogous to the observed situation in the Atlantic, with the added two-layer horizontal transport framework providing additional realism to the current configuration. Because the peak in diazotrophy is collocated with a critical increase in external Fe input, this would be expected to track Fe inputs at steady state. Further, within dynamic model runs incorporating a temporal variability in the region of enhanced Fe input (Fig. S4), i.e., analogous to the latitudinal movement of the ITCZ, the region of enhanced diazotrophy and hence the location of the biogeochemical divide between HPLFe and LPHFe waters, tracks the shifting Fe input in a manner that is qualitatively indistinguishable from the data of our seasonally separated transects (Fig. 3).

Although highly significant biogeochemical responses to seasonal variability in heat input, mixing, and light availability are well known in the open ocean (60, 61), we provide a fully supported description of a large-scale biogeochemical response to seasonal forcing by spatial–temporal variability in external, specifically atmospheric, nutrient inputs. Our results also have important implications beyond the demonstrated seasonal and regional scales. Despite the current lack of comprehensive surface ocean DFe data, the contrasting HPLFe and LPHFe conditions that characterize the South and North Atlantic subtropical gyres (12, 13, 28) appear to be representative of the large-scale biogeochemistry of the entire oligotrophic low bioavailable fixed N (sub)tropical oceans (14, 15, 55). Understanding the processes responsible for this large-scale biogeochemical partitioning is thus of global significance.

Conceptually predictable dynamic responses (Fig. 3) can provide stronger evidence of causation than can be provided by simple correlative studies (12). Consequently, our observations build on those of smaller-event-scale responses of diazotrophs to dust inputs (62, 63). It is crucial to note that the variability in Fe inputs steeply rises to the migration of the ITCZ appears to occur over sufficient spatial and temporal scales that the system is driven fully from the HPLFe to the LPHFe state, over a 300–400 km latitudinal band just north of the equator (Fig. 3). The current study thus potentially represents the closest possible analogy to the type of deliberate large-scale Fe releases that would demonstrate diazotrophy–Fe limitation, in a similar manner to that achieved in the HNLC systems, but which may be practically and politically unfeasible over the long timescales required for fully developed diazotrophic responses (20).

The dynamic response to increased external Fe inputs we describe here thus strongly supports the importance of Fe availability for diazotrophy (1, 11), with consequent excess DIP removal to the point where this subsequently becomes the limiting resource (12, 13, 55). The broad basin-scale biogeochemical partitioning of the oligotrophic oceans into HPLFe and LPHFe regions is thus proposed to be a reciprocal consequence and driver of diazotrophic activity, with enhanced rates of diazotrophy both predicted and observed to occur at the boundaries of these regions (55) (Fig. 3). Continued interpretation of the local, regional, and potentially even the global balance between fixed N loss from the oceans through the processes of denitrification/anaerobic ammonium oxidation and inputs through N₂ fixation (1, 4, 34, 36) will thus need to simultaneously consider spatial patterns of both excess P and Fe availability, as well as variability in these patterns, over multiple timescales. For example, climatic migration of the ITCZ (64) may interact with both regional (38) and remotely forced...
Trace-metal-clean surface seawater samples were collected (7124):163 Trichodesmium and covered an area between 27°N–6°S (9):701. January 28, 2014. We thank the captain and crew of RRS Discovery no. 4 following similar procedures to multifaceted global change both in the past and future. but will also allow better predictions of biogeochemical responses to alterations in natural external forcing, coupled with conceptual mechanistic understanding of the processes involved, not only provides a powerful means for understanding the modern oceans, but will also allow better predictions of biogeochemical responses to multifaceted global change both in the past and future.

Materials and Methods

Water Sampling. Trace-metal-clean surface seawater samples were collected during the UK GA06 GEOTRACES cruise (D361) on the UK research ship RRS Discovery. The cruise took place between February and March 2011 in the (sub)tropical Atlantic Ocean and covered an area between 27°N–7°S and 17°–28°W.

Treatment and On-Board Measurements. All collected DIP samples were analyzed immediately on board with a nanomolar phosphate method using a segmented flow colorimetric analyzer with 2-m liquid waveguide flowcell (69). Dissolved Fe was determined using flow injection analysis (FIA) with luminol chemiluminescence (70). Dissolved Al concentrations were analyzed using FIA with the lumogallion-Al fluorescence technique (72). N₂ fixation was measured using the ¹⁵N₂ technique (72) following similar procedures to those described elsewhere (17, 73).

Numeric Model. The model simulates the interactions between two broad planktonic groups of phytoplankton and diazotrophs in competition for three nutrients (N, P, and Fe) (Fig. S5). The differential equations describing the evolution of the state variables were effectively identical to those used by Ward et al. (55). See the supporting information (SI Materials and Methods) for more detailed information about the model, sample handling, the DFe and DAI techniques, N₂ fixation and associated uncertainties (Fig. S6), and Trichodesmium spp. abundances.

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