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**Observations of
dissolved iron in the
World Ocean**

J. K. Moore and
O. Braucher

Observations of dissolved iron concentrations in the World Ocean: implications and constraints for ocean biogeochemical models

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Abstract

Analysis of a global compilation of dissolved iron observations provides insights into the controlling processes for iron distributions and some constraints for ocean biogeochemical models. The distribution of dissolved iron is consistent with the conceptual model developed for the scavenging of Th isotopes, whereby particle scavenging is a two-step process of scavenging mainly by colloidal and small particulates followed by aggregation and removal on larger sinking particles. Much of the dissolved iron ($<0.4 \mu\text{m}$) is present as small colloids ($>\sim 0.02 \mu\text{m}$) and, thus, likely subject to aggregation and scavenging removal. Only the iron bound to soluble ligands ($<\sim 0.02 \mu\text{m}$) is likely protected from scavenging removal. This implies distinct scavenging regimes for dissolved iron that appear consistent with the observational data: 1) high scavenging regime – where dissolved iron concentrations exceed the concentrations of strongly binding organic ligands; and 2) moderate scavenging regime – where dissolved iron is bound to both colloidal and soluble ligands. The removal rates for dissolved iron will be a function of biological uptake, number and size distributions of the colloidal and small particulate material, ligand dynamics, and the aggregation processes that lead to removal on larger particles.

Inputs from dust deposition and continental sediments are key drivers of dissolved iron distributions. The observations provide several strong constraints for ocean biogeochemical models: 1) similar deep ocean concentrations in the North Atlantic and North Pacific ($\sim 0.6\text{--}0.8 \text{ nM}$), and much lower deep ocean dissolved iron concentrations in the Southern Ocean ($\sim 0.3\text{--}0.4 \text{ nM}$); 2) strong depletion of iron in the upper ocean away from the high dust deposition regions, with significant scavenging removal of dissolved iron below the euphotic zone; and 3) a bimodal distribution in surface waters with peaks less than 0.2 nM and between $0.6\text{--}0.8 \text{ nM}$. We compare the dissolved iron observations with output from the Biogeochemical Elemental Cycling (BEC) ocean model. The model output was in general agreement with the field data ($r=0.76$, for depths $103\text{--}502 \text{ m}$), but at lower iron concentrations ($<0.3 \text{ nM}$) the model is consistently biased high

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relative to the observations.

1 Introduction

Mineral dust supply of dissolved iron to open ocean surface waters was suggested to account for elevated mixed layer concentrations overlying an iron-depleted euphotic zone by Bruland et al. (1994). They estimated a residence time for dissolved iron in the deep ocean of 70–140 years, based on data from the central North Pacific, noting that substantial scavenging removal of iron must occur both in surface waters and in the deep ocean. In a seminal paper, Johnson et al. (1997a) compiled dissolved iron observations from the North Pacific and several additional regions and drew some important conclusions: 1) there are similar concentrations in the deep ocean with no inter-ocean fractionation; 2) iron cycles differently than other highly particle reactive species, likely due to its complexation with organic ligands, which acts to protect iron from removal by scavenging; 3) the continental source for dissolved iron can extend far offshore in the deeper ocean (1000 m) but is removed from surface waters close to shore; 4) dissolved iron concentrations are consistently low in the surface ocean (<0.2 nM). They suggested that the observed iron profiles could be generated by remineralization of a sinking biological particle flux with mean iron/carbon ratio of ~ 5 $\mu\text{mol/mol}$, and particle scavenging removal of iron only where concentrations exceeded ~ 0.6 nM (and the protection of the strong binding iron ligands). They also noted a significant but relatively weak correlation between estimated dust deposition and integrated dissolved iron in the upper 500 m. Most of their observations came from relatively low dust deposition regions in the North Pacific and Southern Ocean.

In the same journal issue there were several comments on the Johnson et al. (1997a) paper. Boyle (1997) suggested that dissolved iron distributions in the deep ocean were likely more varied than implied by the Johnson et al. (1997a) dataset, and that the impact of atmospheric deposition was likely more significant than suggested. These ideas have been supported by subsequent studies showing significantly elevated iron con-

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centrations at the surface and in the deeper ocean beneath the major dust plumes (i.e., Wu and Boyle, 2002; Sedwick et al., 2005), and deep ocean values well below 0.6 nM throughout much of the Southern Ocean (i.e., Measures and Vink, 2001; de Baar et al., 1999; Coale et al., 2005). Sunda (1997) suggested that variations in phytoplankton Fe/C ratios might play a significant role, and estimated the Fe/C ratios in sinking material remineralized in several regions, ranging from $\sim 2 \mu\text{mol/mol}$ in the iron-limited Equatorial Pacific and Southern Ocean regions, to higher values of 7–13 $\mu\text{mol/mol}$ for the high latitude North Atlantic. This analysis assumed minimal subsurface scavenging of dissolved iron; iron removed by such scavenging would imply a higher ratio in regenerated material. In addition, much higher Fe/C ratios in sinking material in the high dust deposition regions would seem necessary to remove excess iron from surface waters. Wu and Boyle (2002) estimated Fe/C export regeneration ratios for the North Atlantic of 23–70 $\mu\text{molFe/molC}$. Lastly, Luther and Wu (1997) suggested iron concentrations will always be a balance between sources and sinks, with organic complexation likely playing a role, and noted that the decrease in surface iron concentrations moving away from the coast was strongly influenced by the width of the shelf due to sediment resuspension events, which released dissolved iron into the water column (see also Chase et al., 2005). In their conceptual model, Johnson et al. (1997a) assumed that there was no particle scavenging of dissolved iron below the concentration of the strong iron binding ligands (0.6 nM), an assumption that has been built into a number of ecosystem/biogeochemical models (i.e., Archer and Johnson, 1999; Lefèvre and Watson, 1999; Aumont et al., 2003). However, in their reply to the comments Johnson et al. (1997b) noted that scavenging would not actually be eliminated at low iron concentrations, as there would always be a small fraction of the iron present as inorganic ions subject to scavenging. Thus, where inputs of dissolved iron were quite low, dissolved concentrations could be reduced below the 0.6 nM value.

In a comprehensive review of iron observations, de Baar and de Jong (2001) noted a significant influence by continental iron sources extending offshore in many regions. They also noted consistently higher iron concentrations ($> 1 \text{ nM}$) in low O_2 regions and

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below the major dust plumes. Low deep water values were noted for the Southern Ocean (~0.3–0.4 nM). They noted that surface values for the open ocean were variable (0.03–0.5 nM) and ranged between 0.3 and 1.4 nM for the deep ocean away from continental influence, with much higher values observed near the coastlines.

5 Martin and coworkers argued that iron was a key limiting nutrient in the oceans, controlling biological production in the High Nitrate, Low Chlorophyll (HNLC) regions of the Southern Ocean, and the subarctic and equatorial Pacific (Martin et al., 1991; Martin, 1992). Subsequent in situ iron fertilization experiments have demonstrated this iron limitation (Coale et al., 1996, 2004; Boyd et al., 2000; Tsuda et al., 2003) and
10 generally confirmed that the entire community is iron-stressed, with the bloom forming diatoms strongly iron-limited, whereas the ambient community, dominated by small phytoplankton, is moderately iron-stressed and experiences strong grazing pressure (Price et al., 1994; see review by de Baar et al., 2005). Model estimates suggest community growth limitation by iron over ~30–50% of the world ocean (Moore et al.,
15 2002b; 2004; Aumont et al., 2003; Dutkiewicz et al., 2005). Observations also suggest that dissolved iron may limit phytoplankton growth rates near the base of the euphotic zone in the subtropical gyres (Bruland et al., 1994; Johnson et al., 1997; Sedwick et al., 2005). In addition, in many subtropical regions the nitrogen fixing diazotrophs may be limited by iron (Falkowski, 1997; Michaels et al., 2001; Berman-Frank et al., 2001;
20 Moore et al., 2004, 2006). This dust-nitrogen fixation linkage may give the subtropics and tropics a sensitivity to atmospheric dust (iron) inputs similar to that seen in the HNLC regions (Michaels et al., 2001; Gruber, 2004). Model estimates indicate that the indirect, nitrogen-fixation-driven biogeochemical response to dust variations can be quantitatively similar to the more direct response in the HNLC regions in terms of
25 total export production and air-sea CO₂ exchange over decadal timescales (Moore et al., 2006). Thus, iron may be the ultimate limiting nutrient for the oceans in the current climate, directly limiting growth in the HNLC regions, and leading to nitrogen being the proximate limiting nutrient in other areas (Moore and Doney, 2007).

Dissolved iron is removed from ocean surface waters through biological uptake and

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through abiotic particle scavenging (adsorption to and removal on sinking particles). Much of our understanding about particle scavenging in the oceans comes from studies of Th isotopes which have a known source through radioactive decay and are not subject to biological uptake. Iron and aluminum are likely scavenged in similar manner, so Th studies have implications for understanding the cycling of these trace metals (Bruland and Lohan, 2004). The conceptual view developed in recent decades is that removal of particle reactive species like ^{234}Th is actually a two-stage process with reversible adsorption, mainly to smaller and colloidal sized particles, followed by aggregation and removal on larger sinking particles (Balistrieri et al., 1981; Bacon and Anderson, 1982; Honeyman et al., 1988; Clegg and Sarmiento, 1989; Honeyman and Santschi, 1989; Wells and Goldberg, 1993; Santschi et al., 2006; see review by Savoye et al., 2006). Models of dissolved Th removal by particle scavenging range from simpler models with a net adsorption rate to sinking particles, to complex models that represent the particle size spectrum down to colloids and explicitly represent the adsorption, desorption, aggregation, and removal processes (Burd et al., 2000; Bruland and Lohan, 2004; see review by Savoye et al., 2006, and references therein).

De Baar and de Jong (2001) noted that the “dissolved” iron measured after passing through a $0.4\ \mu\text{M}$ filter is actually a mix of iron bound to truly dissolved (soluble) ligands ($<0.025\ \mu\text{M}$) and iron bound to fine particulates (colloids $<0.4\ \mu\text{M}$). They suggested a dynamic quasi-equilibrium, shifting iron between organic and inorganic, soluble and colloidal pools, that would be strongly influenced by photochemistry in surface waters, and by the biology that serves as the source for iron binding ligands and drives periodic high export (high scavenging). In the deep ocean, scavenging loss rates would be determined largely by the partitioning between colloidal and dissolved phases, with longer residence times where more Fe was bound by soluble ligands. Wu et al. (2001) studied this division between soluble ($<0.02\ \mu\text{M}$) and colloidal iron ($>0.02\ \mu\text{M}$ and $<0.4\ \mu\text{M}$) using profiles in the subtropical North Pacific and North Atlantic. They found that much of the dissolved iron was present in the colloidal fraction in surface and deep ocean waters, with a colloidal iron minimum in the upper nutricline. They suggested that ag-

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gregation and sinking removal of the colloidal fraction would occur in a manner similar to Th, and that this process should be included in models of oceanic iron cycling. Nishioka et al. (2001) noted considerable temporal and spatial variability in the soluble and colloidal iron concentrations, suggesting a dynamic system. Cullen et al. (2006) built on this work with several profiles in the Atlantic, examining the soluble vs. colloidal fractions of dissolved iron and the ligands that bind iron. They concluded that much of the partitioning of iron between colloidal and soluble pools could be understood by a simple equilibrium partitioning model, but that a significant and varying fraction of the colloidal material seemed inert to ligand exchange with the soluble pool. These studies suggested that the soluble iron may be more bioavailable, as vertical distributions were more like traditional nutrient profiles.

If particle scavenging of iron happens in a similar manner to Th (as seems likely), there are important implications for iron cycling in the oceans. Binding to ligands will not provide complete protection from particle scavenging removal (as often assumed in ocean biogeochemical models), as the colloidal fraction will be subject to aggregation/scavenging removal. Only the truly soluble fraction would be for the most part “protected” from scavenging. The colloid-bound iron would have reduced rates of scavenging loss compared with unbound inorganic iron. This free inorganic iron is a small fraction of the dissolved pool ($< \sim 1\%$), as most dissolved iron is bound to organic ligands (i.e. Rue and Bruland, 1995, 1997; van den Berg, 1995; Wu and Luther, 1995). The particle scavenging removal rate for dissolved iron would then be a function of the proportions in the soluble versus colloidal pools, the dynamics of the particle size distributions and aggregation removal processes, and the turnover/transfer time of dissolved iron between the various forms.

We analyze a global dataset of dissolved iron observations to better understand the sources and sinks for dissolved iron, focusing on open ocean waters and the inputs from atmospheric mineral dust deposition. The Biogeochemical Elemental Cycling (BEC) ocean model simulates the global oceanic distributions of four key phytoplankton functional groups, their potentially growth limiting nutrients (Fe, N, P, and Si), inor-

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ganic carbon, oxygen, and alkalinity (Moore et al., 2004). We compare BEC simulated dissolved iron fields with the available observations, highlighting what the model-data mismatches imply about the underlying model assumptions. In a companion paper (Moore and Braucher this issue, hereafter referred to as MBb) we improve the BEC iron cycle parameterizations and use the model in conjunction with the observations to examine the relative roles of atmospheric mineral dust deposition and continental sediments as sources of dissolved iron to the open ocean.

2 Methods

We analyze field observations of dissolved iron concentrations and compare with simulated iron distributions from the BEC ocean model. The original iron database was compiled by Parekh et al. (2005) and has been expanded by ~30% (to 6540 data points) with data from recent publications. Many of these values are the reported means from duplicate or triplicate samples at a particular depth. Parekh et al. (2005) collected data from the literature and three previous key compilations of iron observations (Johnson et al., 1997; de Baar and de Jong, 2001; Gregg et al., 2003). Some of the data included are dissolved iron concentrations (filter size ranging from 0.2–0.45 μm) and the date, location, and depth of sampling. The complete dataset with references to the original source articles is available as supplementary material. There may be systematic differences in the iron measurements by different groups using different techniques. Ongoing inter-comparison efforts are reducing these differences (Bowie et al., 2003; 2006; and the recent SAFE cruise). Here we make the simplifying assumption that the strong vertical and basin-scale gradients in dissolved iron of interest in this work are larger than these systematic differences.

One focus in this work is evaluating the iron cycle parameterizations in the BEC model for open ocean regions where atmospheric dust deposition is likely the dominant source of dissolved iron. Therefore we have created a subset of the iron database where we attempt to exclude data points strongly influenced by iron coming from other

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sources including the continental margins and shelf/coastal sediments. This is helpful because the remaining subset of data is largely impacted by the mineral dust deposition, and can be used to better understand the associated iron cycling. As a first step we removed data from all ocean model grid cells adjacent to land. This removed much of the observed high iron concentrations associated with shelf/sedimentary sources. However, in some regions it was apparent that the margin influence extended for some distance into the open ocean, particularly at depth, away from the enhanced particle scavenging and biological uptake in the upper ocean.

In Fig. 1, we plot the dissolved iron observations (>1000 m depth) from the eastern subtropical Pacific (latitudes 20–50° N) as a function of approximate distance from the continent. The obvious decline in iron concentrations as distance from the margin increases has been noted previously (Johnson et al., 1997, 2003). The data points within 200 km of the coastline were removed by our land-adjacent rule in the margin-excluded dataset. In addition, we removed all data from the locations in Fig. 1 that are ~550–700 km offshore, retaining the other data points as more representative of the open ocean. There may be influence of the continental margin even more than 1000 km offshore. We address the role of the continental source to the open ocean in our companion paper (MBb, this issue). Through a similar analysis, data was removed several locations near the Asian coast in the NW Pacific. We also exclude data from several papers that measured high iron concentrations attributed to sources not included in the BEC model (riverine or hydrothermal – Mackey et al., 2002; Kerguelen Islands runoff and sediments – Blain et al., 2001; Bucciarelli et al., 2001; and rapid advection from continental sources by the Antarctic Polar Front – Löscher et al., 1997). These steps removed ~48% of the observations, leaving 3176 observations in our open ocean subset (hereafter referred to as the “open ocean” data). This open ocean subset is certainly more impacted by dust and less impacted by sedimentary iron sources than the data that has been excluded. However, in our companion paper we argue that even these open ocean iron concentrations are significantly influenced by the continental sedimentary source for iron (MBb, this issue).

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2.1 BEC model

The coupled biogeochemical elemental cycling (BEC) model (Moore et al., 2002a, 2004) includes ecosystem and biogeochemistry components including full carbonate chemistry dynamics. The model simulates four functional groups of phytoplankton (diatoms, coccolithophores, diazotrophs, and picophytoplankton) and multiple potentially growth limiting nutrients (nitrate, ammonium, phosphate, silicate, and dissolved iron). The BEC runs within the coarse resolution, POP ocean model that is part of the Community Climate System Model (CCSM3.0) developed at the National Center for Atmospheric Research (Collins et al., 2006). The model includes 25 vertical levels with 8 levels in the upper 103 m, a longitudinal resolution of 3.6 degrees and a variable latitudinal resolution, from 1–2 degrees, with the finer resolution near the equator (Collins et al., 2006; Yeager et al., 2006). All the nutrients/elements (C, O, N, P, Si, Fe) are simulated within the full ocean, 3D context with no restoring.

The BEC model reproduces basin-scale patterns of macronutrient distributions, calcification, biogenic silica production, nitrogen fixation, primary and export production (Moore et al., 2002b, 2004). The model has recently been applied to quantify ocean biogeochemical sensitivity to variations in mineral dust deposition (iron inputs) (Moore et al., 2006), the feedbacks between denitrification and nitrogen fixation (Moore and Doney, 2007), and the ocean biogeochemical response to atmospheric deposition of inorganic nitrogen (Krishnamurthy et al., 2007). The results here are from the last year of a 3000 year simulation. This is an extension of the 2000 year control simulation described by Moore and Doney (2007). Iron cycling in the BEC is discussed below, for further details on the BEC model please see Moore et al. (2002a, 2004) and Moore and Doney (2007).

2.2 Iron cycling in the BEC

Dissolved iron sources to the ocean in the BEC model include dissolution of iron from mineral dust particles deposited from the atmosphere and diffusion from shallower

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sediments, while a fraction of the scavenged iron is assumed lost to the sediments to balance these sources (Moore et al., 2004). There is one “dissolved” iron pool that is assumed to be bioavailable, with no distinction between soluble and colloidal forms. A constant fraction of the iron in mineral dust (here 2%) dissolves instantaneously at the surface ocean, with some further iron release through a slower dissolution/disaggregation through the water column (Moore et al., 2004, MBb). Dust deposition is from the climatology of Luo et al. (2003). The sedimentary Fe source is crudely incorporated as a constant flux of $2 \mu\text{mol}/\text{m}^2/\text{day}$ from sediments at the bottom of the ocean grid where depth is less than 1100 m (Moore et al., 2004). As noted by Moore et al. (2004) the coarse resolution ocean grid only weakly captures the bathymetry of the continental shelves. Thus in many coastal/shelf regions the bottom level of the ocean grid is much deeper than the actual shelf regions, providing little iron to surface waters (see MBb, this issue).

Iron is removed from the dissolved pool through biological uptake by the phytoplankton and by particle scavenging. A fraction of the scavenged iron is added to the sinking particulate pool and will remineralize deeper in the ocean, while the remainder is assumed lost to the sediments. Iron scavenging is parameterized in the BEC model based on the mass of scavenging particles and the ambient dissolved iron concentration to crudely account for the presumed influences of iron binding ligands on scavenging losses. Moore et al. (2004) described a scavenging rate that consisted of a base rate times the sinking particle flux divided by a reference particle flux. This approach can be described more simply by combining the base rate and reference flux constant coefficients into one base scavenging rate (F_{base}) that is modified by the sinking particle flux (previously standing stock of POC was also included by Moore et al., 2004). The base scavenging rate ($F_{\text{base}} = 0.01369 \text{ day}^{-1}$) is thus a net adsorption rate to sinking particles, similar to the simplest Th scavenging models. The sinking POC flux and sinking particulate mineral dust fluxes were added to get the particle flux ($\text{ng}/\text{cm}^2/\text{day}$) available to scavenge iron, and a maximum scavenging rate ($\text{MaxPE} = 0.05476 \text{ day}^{-1}$) was imposed (Eq. 1). Given the mean sinking fluxes in the model the

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base scavenging rates including the particle effect at 103 m depth, 502 m, and 2098 m would be $7.5e-4 \text{ day}^{-1}$, $1.6e-4 \text{ day}^{-1}$, and $8.1e-5 \text{ day}^{-1}$, respectively. These rates would be further modified by the ambient iron concentrations. The sinking flux is dominated by POC in surface waters, but due to shorter remineralization length scales for POC, the sinking dust flux becomes important in the deep ocean. POC accounts for 93%, 67%, and 38% of the sinking mass flux at depths of 103 m, 502 m, and 2098 m, respectively.

$$\text{ScavRate} = \text{Febase} * (\text{sinking POC} + \text{sinking Dust}), \text{ must be } \leq \text{MaxPE} \quad (1)$$

$$\text{If } d\text{Fe} > \text{HighFe} \text{ then } \text{ScavRate} = \text{ScavRate} + (d\text{Fe} - \text{HighFe}) * \text{Chigh} \quad (2)$$

$$\text{If } d\text{Fe} < \text{LowFe} \text{ then } \text{ScavRate} = \text{ScavRate} * (d\text{Fe} / \text{LowFe}) \quad (3)$$

$$\text{ScavengedIron} = d\text{Fe} * \text{ScavRate} \quad (4)$$

The scavenging rate increases rapidly at higher iron concentrations (where dissolved iron ($d\text{Fe}$) exceeds HighFe (0.6 nM), $\text{Chigh} = 4286$, Eq. 2) where iron is assumed to begin exceeding the concentrations of strong binding ligands, and progressively decreased at low iron concentrations ($\text{LowFe} = 0.5 \text{ nM}$, Eq. 3) to reflect protection from scavenging losses due to strong iron binding ligands. The scavenging rate is multiplied by the ambient dissolved iron to get the amount removed by scavenging (Eq. 4), of which 10% (F_{Pfe}) is put into the sinking particulate pool, with 90% of scavenged iron presumed lost to the sediments. Most ocean biogeochemical models assume that all scavenged iron is lost from the system (Archer and Johnson, 1999; Christian et al., 2002; Aumont et al., 2003; Parekh et al., 2004, 2005). Gregg et al. (2003) add all scavenged iron to the sinking pool to remineralize at depth, and Doney et al. (2006) send 60% to the sinking particulate pool. Moore et al. (2006) suggested some modifications to the original parameter values for the BEC (similar values are used here see Moore and Doney, 2007).

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3 Results

The observational dataset is heavily weighted towards the upper ocean with 66% of observations from depths less than 103 m, and 86% from depths less than 502 m. The dataset is also weighted strongly towards the Northern Hemisphere (75% of the data).

5 The Southern Hemisphere data is mainly from the Southern Ocean (65% of S.H. data), which we define as latitudes greater than 40.5° S, with few observations in the lower latitudes, mainly in the South Atlantic (18%). There are also strong seasonal biases with only 3.3% of samples collected during winter months. Away from the high deposition regions, particularly at higher latitudes, winter often has maximum surface iron concentrations due to deep mixing and weakened biological uptake. Spring months had the most observations (46%) followed by summer (30%) and fall (21%). Spring is often a time of rapid biological drawdown of dissolved iron. Dust deposition typically has a strong seasonal component peaking during spring or summer months. Thus, iron can have a stronger seasonality, even at low latitudes, that typically seen in oceanographic data.

15 Vertical profiles of dissolved iron tend to follow two patterns: 1) a surface minimum due to surface depletion by biological uptake and scavenging processes; or 2) a surface maximum where there is strong influence by dust deposition events (Johnson et al., 1997a, 2003; de Baar and de Jong, 2001). All observations of dissolved iron are plotted against ocean depth in Fig. 2a, and a similar plot for the open ocean data in Fig. 2b. The symbols used to indicate ocean basin of the samples in Fig. 2 are retained in later figures as well. There are some obvious, strong regional patterns in the iron distributions. The highest surface water concentrations are typically seen in the high dust deposition regions of the northern Indian Ocean and the low latitude North Atlantic (hereafter referred to as the “high deposition regions”), while the lowest surface values are mainly in the Southern and Pacific oceans (Fig. 2). Low surface concentrations are (<0.1 nM) are also seen in some South Atlantic observations. The highest surface concentrations are from coastal waters (with a few points exceeding 10 nM) off of Peru

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(Bruland et al., 2005) and in the Southern Ocean near the Kerguelen Islands (Blain et al., 2001; Bucciarelli et al., 2001). There is considerable variation in surface iron concentrations within basins reflecting differential inputs and removal rates.

In the open ocean dataset, the mean surface iron concentration (≤ 20 m) for all areas except the high deposition regions is 0.25 ± 0.23 nM (where 0.23 is the standard deviation), and the mean in the high deposition regions is 0.76 ± 0.27 nM. The area outside the high deposition regions can be further subdivided between HNLC zones (where annual surface nitrate concentrations exceed $1.0 \mu\text{M}$ in the World Ocean Atlas 2001, Conkright et al., 2001) and the non-HNLC regions. The mean concentration for the HNLC regions is 0.15 ± 0.16 nM and for the non-HNLC regions the mean is 0.27 ± 0.24 nM. It is notable that the iron-limited HNLC regions have mean dissolved iron surface concentrations only moderately lower than the non-HNLC areas. These means are considerably higher than the mean surface value of 0.07 nM calculated by Johnson et al. (1997a). This reflects additional sampling of surface waters shortly after dust deposition events, which in the North Pacific for example can raise surface water concentrations from background levels of < 0.2 nM to values in excess of 0.6 nM (Bruland et al., 1994; Wu et al., 2001; Johnson et al., 2003), increased high latitude sampling early and late in the growing season when deeper mixing may increase iron concentrations, and possibly systematic differences between groups measuring iron.

Many of the elevated deep water concentrations are associated with continental margins and do not appear in our open ocean subset (compare Figs. 2a and b) most notably in the Pacific data. Deep water values in the open ocean dataset range mainly between ~ 0.2 – 1.0 nM, with the Southern Ocean showing consistently low values and the North Atlantic and North Pacific having higher, similar concentrations. Unlike the North Atlantic data which typically have a surface maximum and decline with depth, the Arabian Sea data tend to increase modestly with depth in the upper ocean (Fig. 3). This may be partially due to lower scavenging losses in the sub-oxic waters below the euphotic zone (Measures and Vink, 1999).

There is a clear signal from dust deposition in the depth-resolved iron data with the

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5 highest concentrations in the open ocean dataset all in the high deposition regions, and with the lowest upper ocean iron concentrations in the low deposition areas of the Southern Ocean and the equatorial Pacific. In Fig. 3, we plot all dissolved iron observations from the open ocean dataset against the climatological annual dust deposition estimated by Luo et al. (2003) for the late 20th century. There is considerable uncertainty in these model estimates of dust deposition, but some general patterns seem fairly robust. Dust deposition varies over three orders of magnitude while most of the iron observations fall within a much narrower range (~two orders of magnitude). This reflects the non-linear nature of iron removal processes, although variations in aerosol iron solubility may also play a role, as the lower deposition areas generally farther from source regions may have higher solubilities (see Mahowald et al., 2005, and references therein). The regions beneath the major dust plumes in the North Atlantic and northern Indian oceans receive two orders of magnitude higher dust deposition than most other areas. The North Pacific receives ~2–5 times more dust than most Southern Ocean sites, although some of the very lowest deposition rates are in the equatorial Pacific. There is some overlap across all the regions, some North Atlantic sites receive low dust levels, and the South Atlantic sites span the range from low to high deposition. A few Southern Ocean regions receive more than 1 g dust/m²/yr (Fig. 3). For each dust deposition rate there is typically a wide range of observed iron concentrations. This likely reflects errors in the dust deposition, variable removal by biological uptake and particle scavenging, influence by other Fe sources, and variations in depth of the sample (although the database is strongly weighted towards the upper ocean).

20 There was sufficient data in the North Atlantic, North Pacific, and Southern Ocean to calculate mean profiles of dissolved iron from the open ocean dataset (Fig. 4). In the upper ocean mean dissolved iron concentrations were quite low in the Southern Ocean (0.16 nM upper 100 m) and in the North Pacific (0.20 nM upper 100 m), with much higher iron levels in the North Atlantic (0.73 nM). There was a sub-surface minimum in the North Atlantic between 100–250 m depth, noted previously (Wu et al., 2001; Wu and Boyle, 2001; Sedwick et al., 2005; Bergquist and Boyle, 2006). Below

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250 m the profiles for the North Atlantic and North Pacific basins are similar, despite a roughly two orders of magnitude difference in dust deposition (Figs. 3 and 4). This similarity was noted previously in a much smaller dataset by Johnson et al. (1997a). The Southern Ocean has much lower mean concentrations below 250 m with the largest difference between 1000–1500 m by a factor of ~ 2.6 – 2.9 . Values appear to converge somewhat in the deepest ocean, but there were few observations below 2000 m (see Fig. 2b). We calculated mean iron concentrations below 500 m depth for these basins as 0.37 ± 0.19 (n=96) for the Southern Ocean, 0.74 ± 0.33 nM (n=107) for the North Atlantic, and 0.74 ± 0.21 nM (n=232) for the North Pacific. The North Pacific data may reflect a strong influence from continental sources (see Fig. 1 and MBb). If all observations are included (rather than just the open ocean subset) mean concentrations are higher in the North Pacific (0.87 ± 0.37 nM, n=468) than in the North Atlantic (0.76 ± 0.31 nM, n=149), reflecting in part more sampling near the continents in the North Pacific. The Southern Ocean mean for all data was 0.46 ± 0.28 nM (n=160).

The BEC simulated iron distributions are compared with the open ocean observations in Fig. 5 (with model values extracted from the same month, depth, and location as the observations). There is a broad agreement between the model and the observations in surface waters (r-correlation coefficient of 0.73, Fig. 5a). However, the model tends to overestimate iron concentrations, particularly at the lower iron concentrations, below ~ 0.3 nM. Values at the high end of iron concentrations also tend to be somewhat too high in the model. There are a couple of dozen points in the Atlantic basin where the model drastically overestimates iron concentrations (Fig. 5a). This reflects the coarse resolution of the ocean model and the atmospheric model used to generate the dust deposition field. Due to the sharp gradients involved small shifts in the location of the dust plume can create large errors in the model-data comparison, leading to the model estimating high iron concentrations at a number of points where low iron was observed. In subsurface waters the model again is relatively well correlated with the observations (r=0.76, 0.70 for log-transformed data, Fig. 5b). In contrast to surface waters the model underestimates concentrations at the high end (>0.6 nM) and again

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tends to overestimate iron concentrations at the low end ($< \sim 0.3\text{--}4\text{ nM}$) (Fig. 5b). There is significant seasonal variability in the surface and subsurface iron concentrations in the high deposition regions (not shown).

We compare spatial plots of annual mean BEC simulated iron with the observations, illustrating the open ocean subset and full dataset observations (Figs. 6–8). Looking at the surface water observational data (0–103 m) dissolved iron concentrations are typically quite low ($< 0.2\text{ nM}$) outside the high deposition regions (Fig. 6). The model displays qualitatively similar patterns, but often overestimates the absolute concentrations in the low iron regions. In part this is due to seasonal entrainment of iron at higher latitudes (and observational under-sampling during winter months), but the model estimates were too high even when comparing with monthly model output from the same months as the observations (Fig. 5).

In subsurface waters the tendency for the model to overestimate iron concentrations away from the high deposition regions is even more apparent (Fig. 7). Given our inputs of dissolved iron from dust deposition, it is apparent that strong scavenging removal of iron in the upper water column below the euphotic zone is necessary to reproduce the observed concentrations. The model tends to underestimate the subsurface dissolved iron concentrations in the high deposition regions (Fig. 7). In the deep ocean ($> 500\text{ m}$) the simulated iron concentrations are lowest in the eastern Equatorial Pacific followed by the Southern Ocean, with high concentrations ($> 0.6\text{ nM}$) in the high deposition regions (Fig. 8a). The model output in the deep North Pacific has concentrations in between the North Atlantic and Southern Ocean regions, a pattern suggested by Parekh et al. (2005) for the real ocean. However, the observations don't strongly display this pattern, but have values in the North Pacific that are similar to the high deposition regions, except for the data closest to the central portion of the gyre where iron concentrations are lower. Examining the full dataset in this region it appears that coastal iron sources extend influence well into the North Pacific gyre, partially compensating for the lower dust inputs from the atmosphere (Figs. 1 and 8c, see also MBb). The simulated iron concentrations in the deep Southern Ocean are considerably higher than

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observations. The model only weakly captures the strong contrast between deeper iron concentrations in the North Pacific and Southern Ocean (Figs. 7 and 8).

In the surface observations from the Ross Sea and off the western coast of North and South America, high concentrations are apparent in the observations near land, declining quickly offshore (Fig. 6c). A similar pattern can be seen in the subsurface observations in the Ross Sea, the eastern North Atlantic, the eastern North Pacific, the gulf of Alaska extending southwards from land, and in the western North Pacific (Fig. 7c). At these subsurface depths (103–502 m) the coastal source can often be seen to extend farther offshore, than in surface waters (Figs. 6c and 7c). This pattern extends into the deep ocean as well (Fig. 1 and Fig. 8c, see also MBb) due to reduced particle flux and scavenging losses at depth.

4 Discussion

We have seen strong regional to basin scale trends in the observed distributions of dissolved iron in the world ocean. These patterns confirm the strong role of atmospheric dust deposition as a source of dissolved iron to the open ocean (Figs. 2 and 3). However, this dust-iron relationship is highly nonlinear (Fig. 3), reflecting the strong biological influence on iron distributions. Mean deep ocean iron concentrations are similar in the North Pacific and North Atlantic, despite the much higher dust inputs to the Atlantic (Fig. 4). It appears that the coastal iron source may contribute substantially to the open ocean in the North Pacific, partially compensating for the lower dust inputs. We address the relative roles of atmospheric and sedimentary iron sources in the companion paper (MBb, this issue).

Despite a wide range of dust inputs from the atmosphere, the surface iron distributions exhibit a strong bi-modal distribution, with peaks at ~ 0.1 – 0.2 nM and at ~ 0.6 – 0.8 nM. These patterns likely result from interactions with the organic ligands that bind nearly all the dissolved iron in seawater. These interactions appear to set up distinct scavenging regimes for dissolved iron in the oceans, depending on the balance

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between iron sources and sinks. At very high iron inputs, as in the high deposition regions, the strong binding ligands can become saturated, and scavenging rates will increase as an increasing proportion of iron is bound to weaker ligands or exists as free inorganic iron. Some removal of the colloidal size class of ligands will occur due to aggregation and settling processes, but this will rapidly be replenished in surface waters, where biological processes releasing the ligands are active, and perhaps in the deeper ocean during remineralization of sinking organic matter. At lower iron input levels, this aggregation and removal of the colloidal size class will decrease surface dissolved iron concentrations to lower concentrations. Biological uptake will also remove substantial amounts of dissolved iron, particularly as some larger phytoplankton engage in luxury uptake of iron. At very low iron input levels, or as these processes deplete iron down to very low levels ($<0.1\text{--}0.15\text{ nM}$), the loss rate for dissolved iron will likely decline. A higher proportion of the ligand-bound iron may exist in the soluble size class (Nishioka et al., 2001, 2005) and, therefore, will not be subject to significant aggregation/scavenging removal. Also, the phytoplankton uptake will decrease as available iron approaches the half-saturation values for iron uptake and phytoplankton become increasingly iron stressed (growing more slowly and decreasing their cellular Fe/C ratios). Thus, outside the high iron input areas, dissolved iron concentrations will tend to be depleted to low levels in surface waters ($<0.2\text{ nM}$), over a fairly wide range of iron inputs from mineral dust and other sources.

Our assumption in the BEC model that scavenging rates progressively decrease as iron decreases below 0.5 nM is incorrect and leads to an overestimation of dissolved iron concentrations at the low end of observations (Figs. 5–9). In the companion paper, we demonstrate a much better BEC fit to observations using a first-order, particle-dependent scavenging rate, with no dependence on ambient iron concentration for iron values below 0.6 nM (MBb, this issue). Our model results also suggest that there must be significant removal of dissolved iron in subsurface waters, where the iron concentrations are typically well below 0.6 nM , for the model to match the observed iron distributions (MBb, this issue). Thus, aggregation and scavenging removal of the col-

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loidal iron pool likely plays a key role in determining dissolved iron concentrations (Wu et al., 2001; de Baar and de Jong, 2001; Nishioka et al., 2001).

At present there are only a few studies of the size-fractionation of “dissolved” iron between the soluble and colloidal pools. The soluble pools have been noted to have strongly nutrient type profiles, while the colloidal fraction is more variable, often with a subsurface/upper thermocline minimum (Wu et al., 2001). Soluble Fe concentrations in surface waters have been measured at ~ 0.1 nM in oligotrophic (presumably N or P limited) regions (Wu et al., 2001; Wen et al., 2006), although Cullen et al. (2006) found values ranging between 0.04–0.28 nM in the Atlantic. Somewhat lower values of ~ 0.05 nM have been reported in the iron-limited subarctic NE Pacific and Southern Ocean (Nishioka et al., 2001; 2005). In these studies, the concentration of soluble iron increased steadily with depth, to values ~ 0.2 – 0.4 nM below several hundred meters.

The increasing soluble iron concentrations with depth may partially explain the observed trend towards increasing iron concentrations with increasing depth (Figs. 6–8, see also Fig. 4 in MBb). Other factors also likely contribute, including strong biological drawdown in surface waters, and weaker scavenging in the deep ocean due to lower particle concentrations. Aggregation and scavenging of the colloidal iron fraction may also explain the observed subsurface minimum concentration for colloidal iron (Wu et al., 2001). In these relatively shallow, subsurface waters, iron inputs and likely the production of the organic ligands are lower than in surface waters, but particle flux, colloidal aggregation, and scavenging rates are likely fairly high. Thus, even in high deposition regions, much of the colloidal fraction may be removed by aggregation and scavenging. The release of this scavenged iron, along with lower particle flux and scavenging rates, then leads to higher iron concentrations in the deep ocean. Scavenging rates would still increase as dissolved iron rose above ~ 0.6 – 0.7 nM, setting an upper limit on the deep ocean concentrations (Fig. 8c), as suggested by Johnson et al. (1997a). At any one location, the observed profiles are likely fairly dynamic and dependent on the balance between the various local sources and sinks and advective processes (Wu and Luther, 1997; Wu and Boyle, 2002). Nishioka et al. (2001) noted strong temporal

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variations in the soluble iron fraction at depth. There is likely a dynamic, rapid turnover and exchange of iron amongst the various pools, both at the surface and in the deeper ocean (de Baar and de Jong, 2001; Bruland and Lohan, 2004; Cullen et al., 2006).

A global collection of dissolved iron observations has now accumulated that is sufficient to place some strong constraints on ocean biogeochemical models. There appears to be substantial scavenging removal of iron in areas where ambient concentrations are well below 0.6 nM. Thus, models that assume scavenging removal only when iron concentrations exceed this threshold will tend to overestimate subsurface iron concentrations and subsurface Fe inputs to the euphotic zone. This will tend to over-emphasize the subsurface iron source and underestimate the importance of dust deposition as a driver of ocean biogeochemical cycling. Thus, it is critical that models be evaluated against observations both for surface fields, and for the subsurface depths that drive much of the iron input to surface waters (see also MBb).

In the high deposition regions, observed iron concentrations are often well above 0.6 nM, although this signal appears to be removed by scavenging over relatively short timescales (weeks to months), allowing for limited advection. The mean surface iron concentration for areas outside the high deposition regions is 0.25 ± 0.23 nM, and the mean in the high deposition regions is 0.76 ± 0.27 nM. This is only a three-fold difference in mean iron concentrations, despite variations in iron inputs of several orders of magnitude. The mean concentration for the HNLC regions is 0.15 ± 0.16 nM, only modestly lower than for the non-HNLC regions (excluding high deposition areas) where the mean is 0.27 ± 0.24 nM. These averages include sporadic increases due to dust deposition events (i.e., Bruland et al., 1994; Johnson et al., 2003), otherwise surface values are typically below 0.2 nM. The deep North Pacific has similar deep ocean concentrations to the high deposition regions, with strong lateral gradients – high iron values near the continental margins, decreasing towards the center of the basin. Another key metric is how well models can reproduce the binned distributions of iron observations (MBb, this issue).

Hard work by numerous researchers has slowly built up the current global iron

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database over several decades, but there is still much we do not know about the marine iron cycle. The observational database for iron is still a tiny fraction of that available for other key nutrients. The size-fractionation of the dissolved pool and the relative roles of soluble and colloidal classes are only just beginning to be addressed. Numerous questions remain concerning the particle scavenging, bioavailability, photochemistry, and turnover times of the various iron pools. The iron fertilization experiments have demonstrated the key role iron plays in governing ocean productivity and biogeochemical cycling (de Baar et al., 2005). Fully unraveling all the mechanisms and processes controlling iron cycling in the modern ocean, and incorporating these results into ocean biogeochemical models to understand the role of iron cycling in future and past climate regimes, will be a key area of research in the coming decades.

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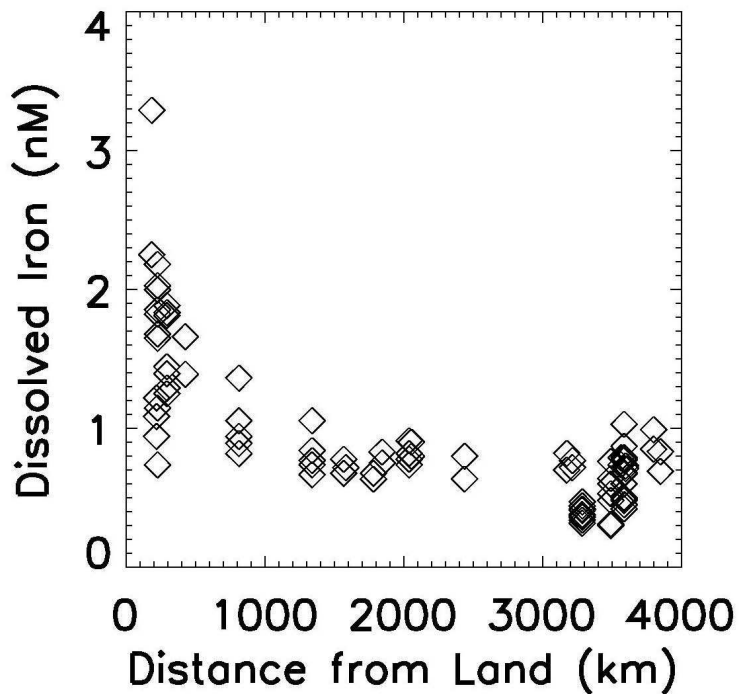


Fig. 1. Observed iron concentrations from depths greater than 1000 m in the eastern subtropical Pacific Ocean (20–50° N) plotted as a function of distance to the continental land mass.

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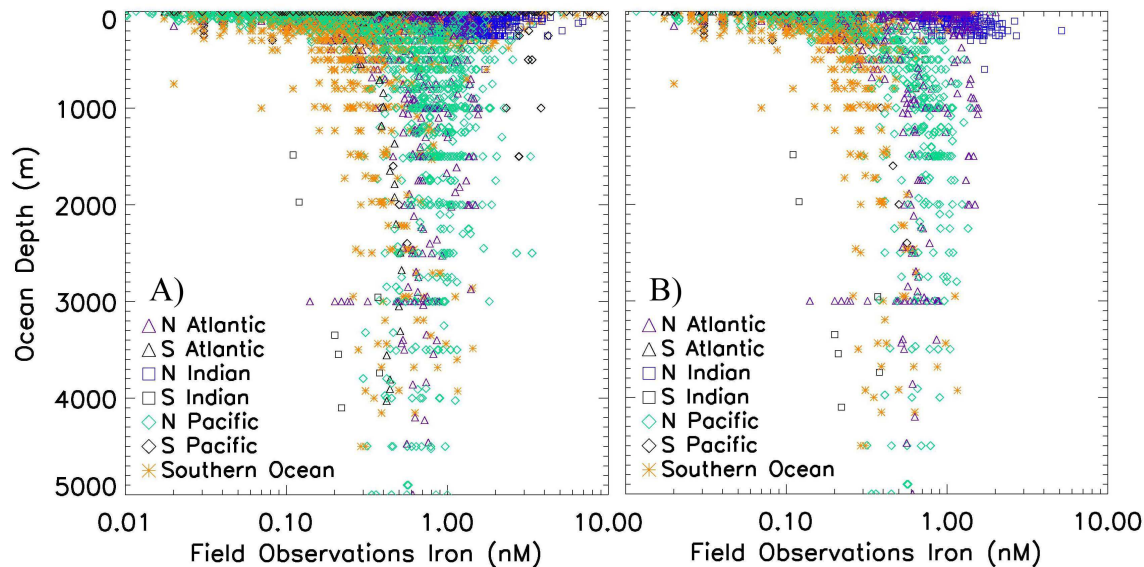


Fig. 2. All observations of dissolved iron plotted as a function of depth with symbols denoting ocean basins (A); all observations from the “open ocean” subset (see text for details) plotted as a function of depth with symbols denoting ocean basin.

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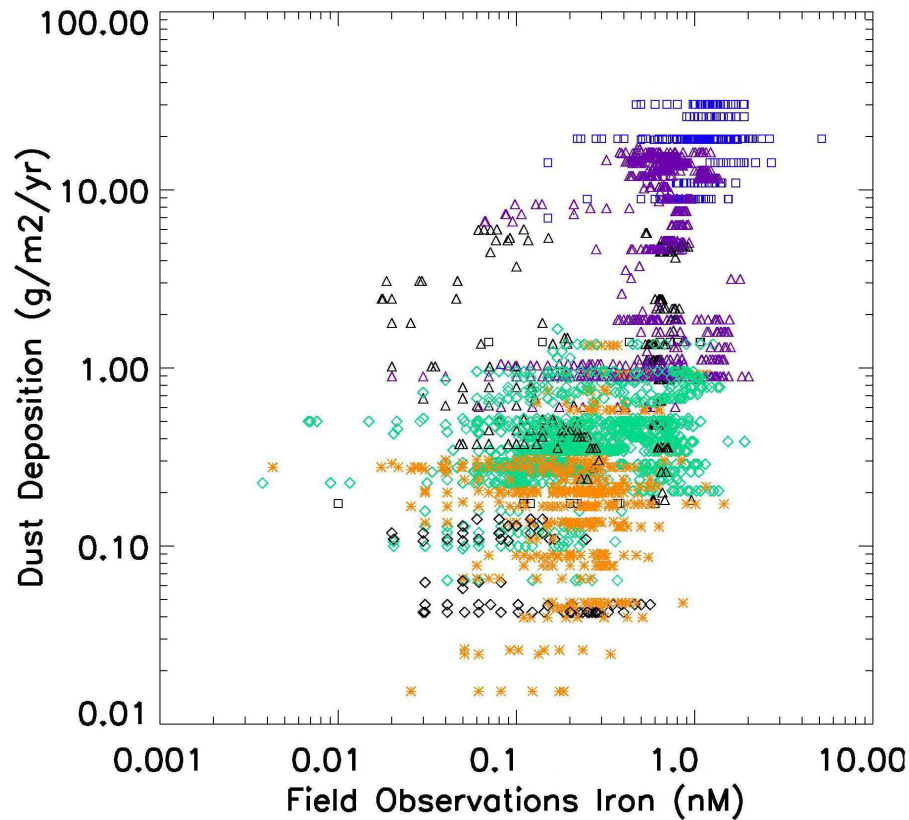


Fig. 3. All observations of dissolved iron from the open ocean subset (see text for details) plotted against atmospheric transport model estimates of annual mineral dust deposition to the oceans in the climatology of Luo et al. (2003). Symbols indicate ocean basin as in Fig. 2.

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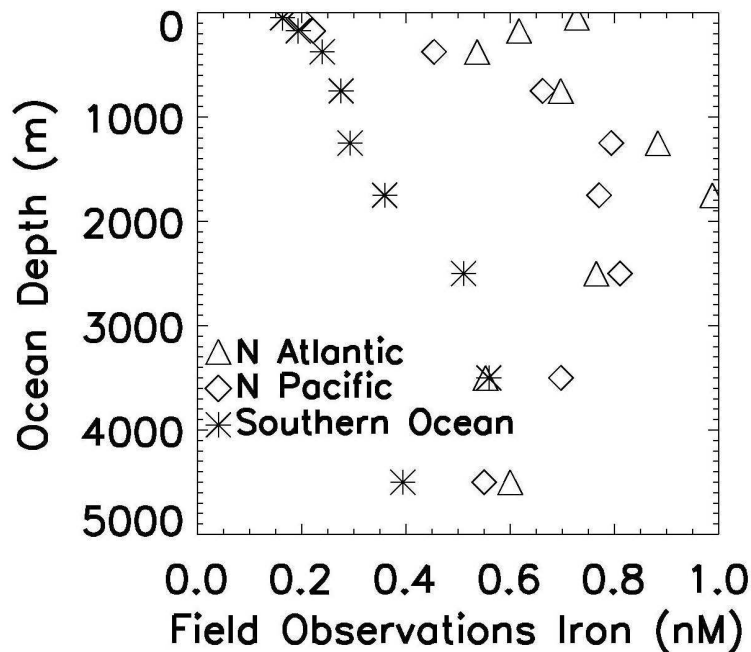


Fig. 4. Mean profiles of dissolved iron in the North Atlantic, North Pacific, and Southern Ocean averaged over the depth intervals: 0–100 m, 100–250 m, 250–500 m, 500–1000 m, 1000–1500 m, 1500–2000 m, 2000–3000 m, 3000–4000 m, 4000–5000 m.

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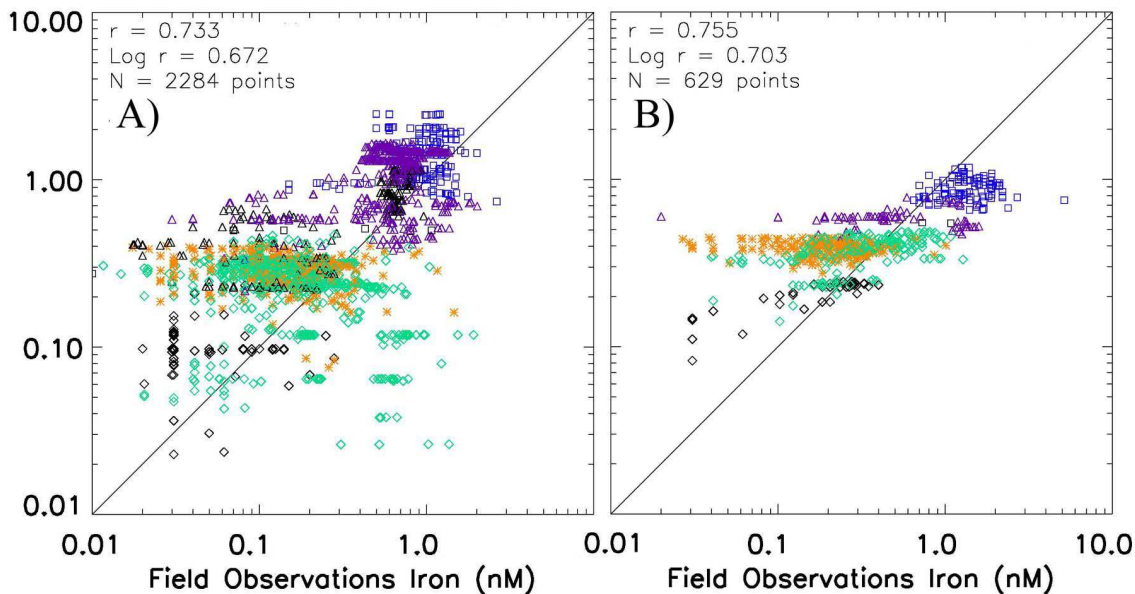


Fig. 5. Iron observations from the upper 103 m plotted against BEC simulated iron from the same month, depth, and location (A); same plot but for depths 103–502 m (B).

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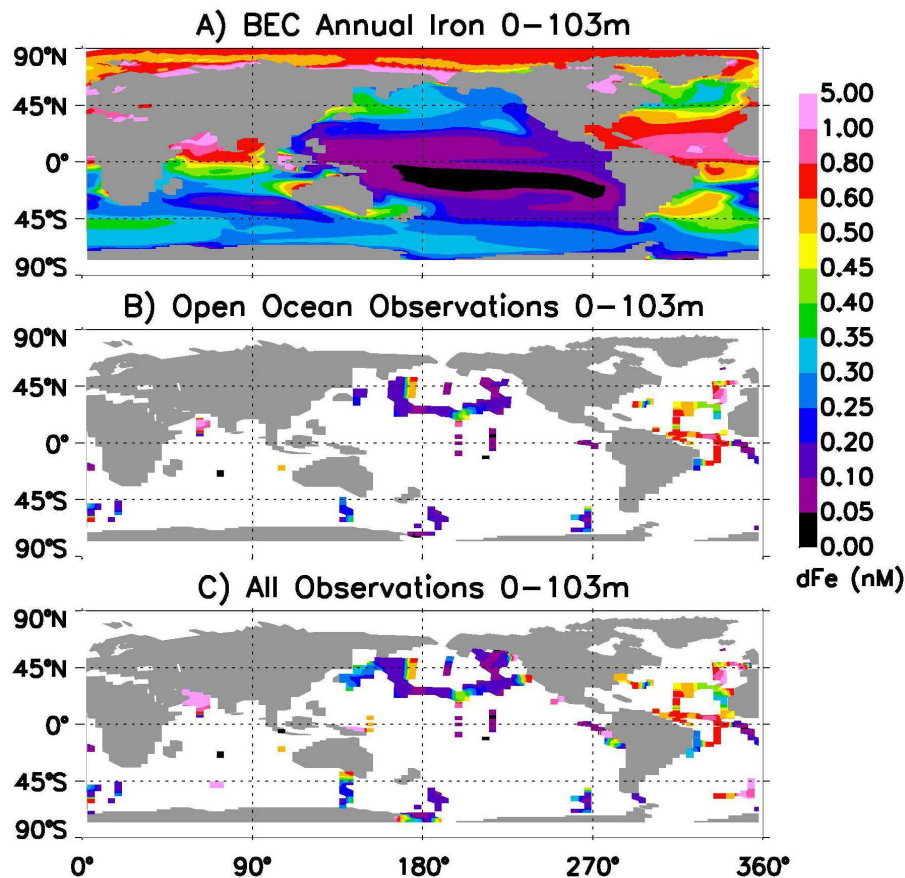


Fig. 6. Iron observations averaged onto CCSM3 ocean grid from depths less than 103 m compared with annual mean BEC simulated iron concentrations averaged over the upper 103 m.

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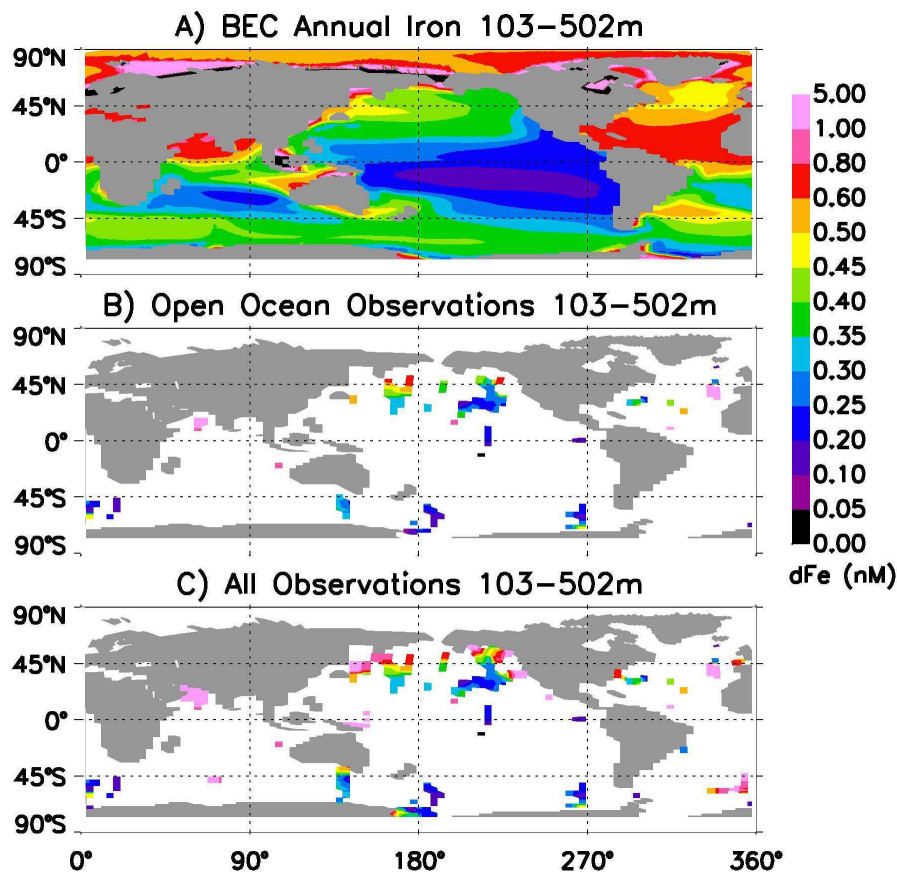


Fig. 7. Iron observations averaged onto the CCSM3 ocean grid from depths greater than 103 m and less than 502 m compared with annual mean BEC simulated iron concentrations averaged over this depth range.

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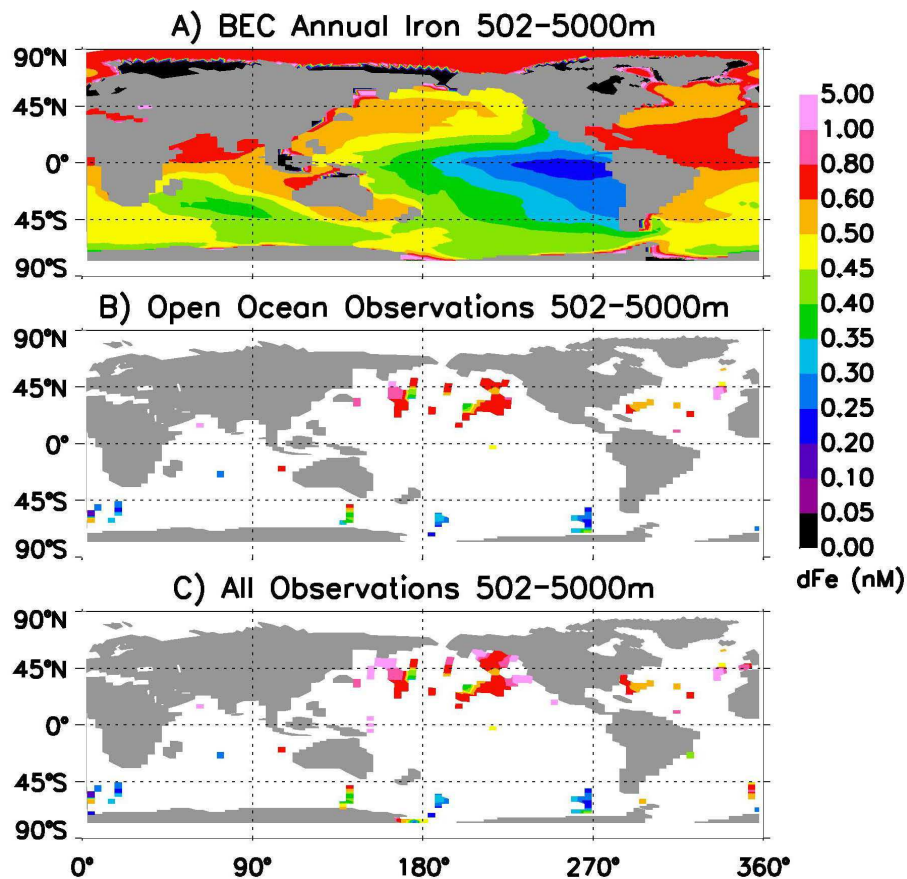


Fig. 8. Iron observations averaged onto the CCS3 ocean grid from depths greater than 502 m compared with annual mean BEC simulated iron concentrations averaged over depths below 502 m.

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