

Supplementary information for: Manganese co-limitation of phytoplankton growth and major nutrient drawdown in the Southern Ocean

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Supplementary Tables

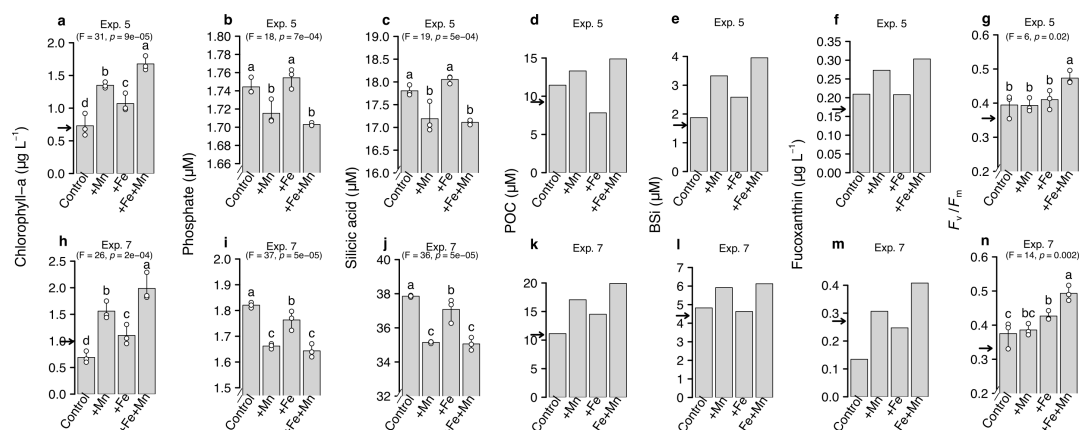
Supplementary Table 1. Experiment starting conditions.

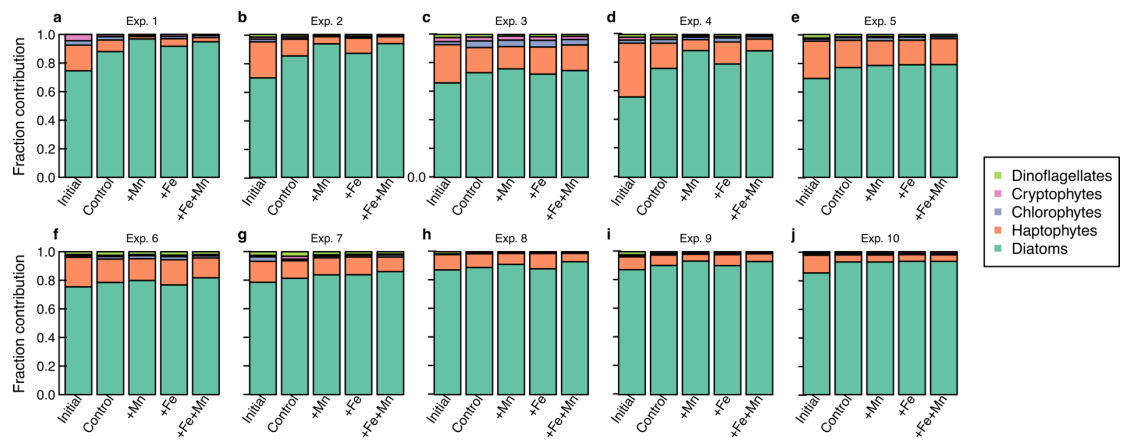
Exp.	Lon. (°E)	Lat. (°N)	Duration (Days)	SST (°C)	$E_{in situ}/E_{exp}^{\dagger}$	NO_3^- (μM)	PO_4^{3-} (μM)	$Si(OH)_4$ (μM)	DFe (nM)	DMn (nM)	Mn* ^(‡) (nM)	Chl.-a (mg m ⁻³)
1	-58.01	-54.71	3.1	5.2	1.01	21.07	1.44	6.79	0.35	0.29	0.16	0.82
2	-57.65	-55.44	2.9	5.4	0.57	20.53	1.39	6.39	0.16	0.37	0.31	1.59
3	-57.96	-55.60	1.8	5.1	1.03	21.34	1.45	6.71	0.39	0.36	0.21	0.75
4	-57.79	-55.76	2.9	5.2	0.72	21.49	1.49	7.57	0.25	0.33	0.24	0.91
5	-57.38	-56.55	3.2	2.2	0.66	27.41	1.75	18.85	0.44	0.15	-0.02	0.65
6	-57.23	-56.81	3.2	1.6	0.46	27.38	1.77	20.05	0.39	0.14	-0.01	0.59
7	-56.42	-58.08	5.1	-0.4	1.01	27.31	1.79	36.12	0.34	0.17	0.04	1.21
8	-56.09	-58.66	3.7	-0.2	0.52	26.32	1.57	41.81	0.52	0.38	0.19	2.18
9	-55.47	-59.64	3.1	-0.1	0.97	26.68	1.59	46.40	0.42	0.34	0.18	2.97
10	-54.64	-60.97	3.9	-0.6	1.99	29.83	2.08	78.52	0.78	2.35	2.06	1.26

[†]Ratio of estimated in situ mean mixed layer irradiance to mean experimental irradiance

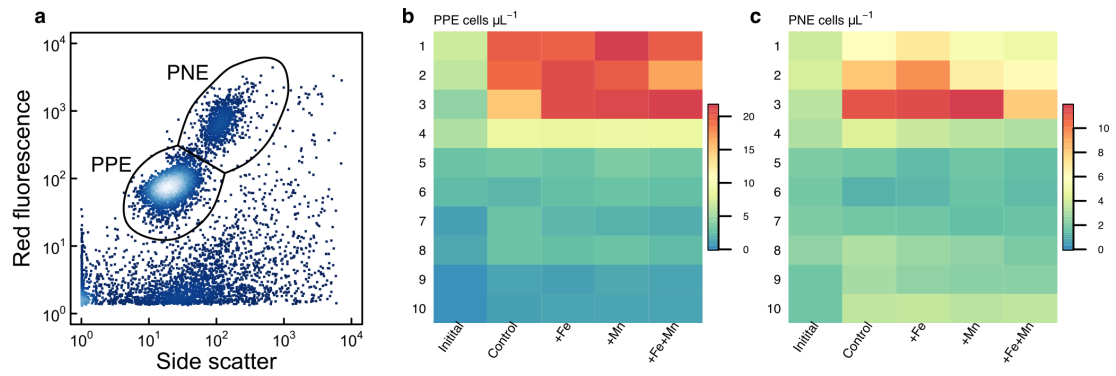
[‡]Mn*=DMn – DFe/ $R_{Fe:Mn}$, where $R_{Fe:Mn}$ represents the assumed-average Fe:Mn ratio of phytoplankton (2.67)¹.

Supplementary Figures

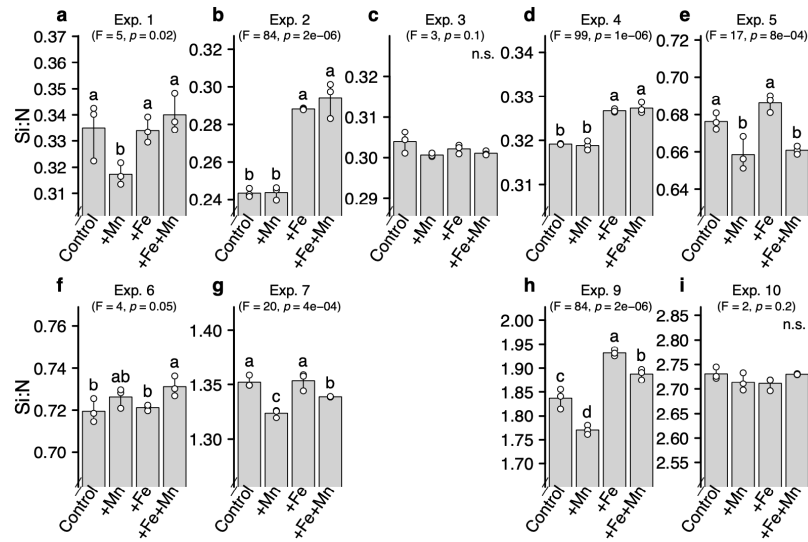




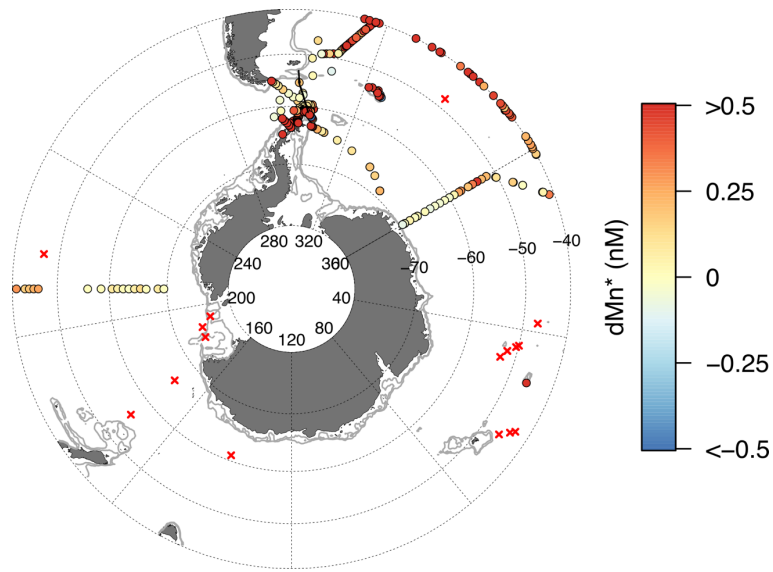
Supplementary Figure 2. Phytoplankton community composition. Predicted contribution of different phytoplankton types to total chlorophyll-a via diagnostic pigment analysis and CHEMTAX.



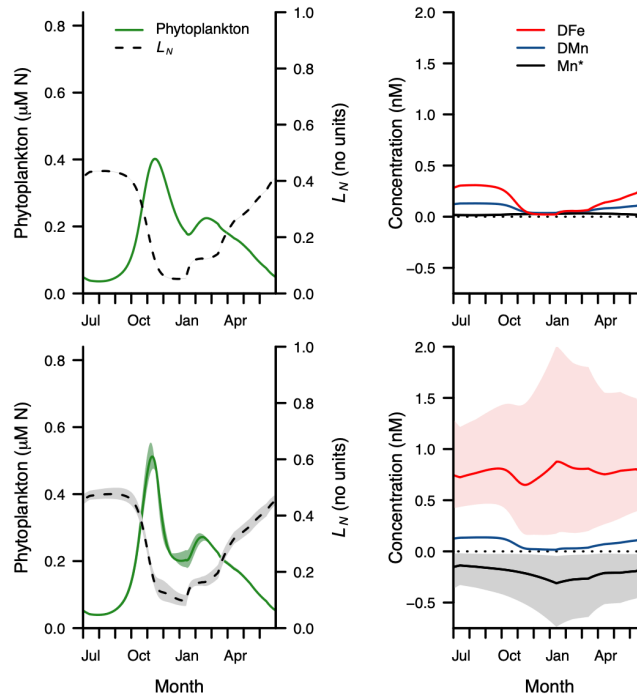
Supplementary Figure 3. Responses of small phytoplankton analyzed by flow cytometry. **a**, Flow cytometry gating strategy. PPE indicates photosynthetic picoeukaryotes (approximately 0.2–2 μm); PNE indicates photosynthetic nanoeukaryotes (approximately 2–20 μm). Gate positioning was checked and adjusted manually for each sample to account for changes in cellular fluorescence. The units for the axes are arbitrary. **b**, photosynthetic picoeukaryotes. **c**, photosynthetic nanoeukaryotes.



Supplementary Figure 4. Changes in dissolved silicic acid: nitrate concentration ratios (Si:N) in experiments. Bar heights indicate the mean, white symbols are the individual replicate values (connecting line is the range), and different letters above bars indicate statistically different mean responses (n=3 biological replicates; one-way ANOVA, F and p values shown, followed by Fisher Least Significant Difference test; n.s. indicates not significant). Results for Experiment 8 are not shown as samples were stored frozen for >1 year prior to analysis (see Methods; they however showed significant silicic acid: nitrate drawdown in +Fe and +Fe+Mn additions relative to controls and +Mn).



Supplementary Figure 5. Wider extent of Mn deficiency in the surface Southern Ocean. Values are calculated with <30 m depth data from the GEOTRACES Intermediate Data Product², CLIVAR^{3,4}, and Refs 5–11. Yellow points indicate sites approaching Mn-Fe co-deficiency. Red crosses indicate locations where experiments (replicated bottle-scale bioassay experiments or mesoscale enrichment experiments) have found evidence for Fe limitation (from compilation of Ref. 1).



Supplementary Figure 6. Ecosystem model simulation in response to enhanced dust. Details are as for Figure 3d–g and Methods, except (i) all panels are for sites with low open ocean Mn*, isolated deep-water case; and (ii) dust inputs have been included in simulations. Upper panels are for interglacial, low dust conditions ($0.014 \text{ g m}^{-2} \text{ yr}^{-1}$)¹²; lower panels are for glacial high dust conditions ($3.45 \text{ g m}^{-2} \text{ yr}^{-1}$)¹³. The Fe and Mn content of dust were prescribed (Fe=30,890 p.p.m.; Mn=527 p.p.m.)¹⁴. Solid lines represent results when solubilities of Fe and Mn are set to the medium values found in the compilation of Ref. 15, whilst shading represents the range when lower and upper quartile values of the compilation are used (see main text and Methods).

Supplementary References

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