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Supplement of

Increasing P limitation and viral infection impact lipid remodeling of the picophytoplankter *Micromonas pusilla*

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1 Table S1: Averages \pm standard deviations of relative IPL peak areas under the different P-
2 treatments. Replicates (n=2) are derived by pooling the high and low pCO₂ data which didn't
3 show a significant difference ($0.071 < p < 0.623$).

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	SQDGs	PGs	DGDGs	MGDGs	DGTSs	DGTAs
	(%)	(%)	(%)	(%)	(%)	(%)
P-replete	3.0 \pm 0.1	10.5 \pm 0.7	2.8 \pm 0.1	9.5 \pm 0.8	2.7 \pm 0.6	71.5 \pm 0.7
0.97 μ_{\max} P-controlled	8.8 \pm 0.5	6.0 \pm 1.9	7.5 \pm 0.4	5.7 \pm 0.6	4.5 \pm 1.4	67.5 \pm 2.1
0.32 μ_{\max} P-controlled	8.9 \pm 0.2	4.5 \pm 0.0	9.5 \pm 0.1	6.5 \pm 0.7	2.3 \pm 0.2	68.5 \pm 0.7

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7 Figure S1: Relation of the SQDG:PG (A) and DGDG:PG (B) ratios with alkaline phosphatase
8 activity (APA) of the P-replete (circles) and $0.97\mu_{\max}$ (squares) and $0.32\mu_{\max}$ (triangles) P-
9 controlled cultures. The ratios of the virally infected cultures are depicted in grey. The
10 culturing treatments are not depicted in the culture but can be derived from Table 1 (APA
11 value for each treatment). Linear regressions for both ratios were significant for the $0.97\mu_{\max}$
12 cultures ($n=5$, $p<0.01$, $r^2=0.99$; normality by Shapiro-Wilk test: $p=0.258$ and 0.378 for the
13 SQDG:PG and DGDG:PG ratio, respectively).

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