Lipidomics of *Thalassiosira pseudonana* Under Phosphorus Stress Reveal Underlying Phospholipid Substitution Dynamics and Novel Diglycosylceramide Substitutes

Running Title: P-Stress Diatom Lipidomics

Jonathan E. Hunter\(^1,2\)*, Joost Brandsma\(^3\), Marcus K. Dymond\(^4\), Grielo Koster\(^3\), C. Mark Moore\(^1\), Anthony D. Postle\(^3\), Rachel A. Mills\(^1\) and George S. Attard\(^5\).

1. Ocean and Earth Science, University of Southampton, National Oceanography Centre, Southampton, European Way, Southampton, SO14 3ZH, United Kingdom
2. Institute for Life Sciences, University of Southampton, SO17 1BJ, United Kingdom
3. Faculty of Medicine, University of Southampton, Southampton General Hospital, Tremona Road, Southampton, SO16 6YD, United Kingdom
4. Division of Chemistry, School of Pharmacy and Biomolecular Sciences, University of Brighton, Brighton, BN2 4GJ, United Kingdom
5. Chemistry, University of Southampton, Southampton, SO17 1BJ, United Kingdom

†Corresponding Author: jhunter@whoi.edu; Current Address: Marine Chemistry and Geochemistry, Woods Hole Oceanographic Institution, Woods Hole, MA 02543-1050, United States of America
Abstract

Phytoplankton substitute phosphorus (P) containing lipids with non-P analogues boosting growth in P-limited oceans. In the model diatom *Thalassiosira pseudonana*, lipid headgroup substitution dynamics are well described, but those of the individual lipids, varying in fatty acids, are unknown. Moreover, the behaviour of lipids outside of the common headgroup classes, and the relationship between lipid substitution and cellular particulate organic P (POP) are yet to be reported. We investigated these through the mass spectrometric lipidomics of P replete (P+) and depleted (P-) *T. pseudonana* cultures. Non-lipidic POP was depleted rapidly by the initiation of P stress, followed thereafter by cessation of P-lipid biosynthesis and per-cell reduction in P-lipid of successive generations. Minor P-lipid degradative breakdown was observed, releasing P for other processes, but most remained intact. This may confer an advantage to efficient heterotrophic lipid consumers in P limited oceans. Glycerophosphatidylcholine (PC), the predominant P-lipid, was similar in composition to its betaine substitute lipid. During substitution, PC per cell was less abundant and more highly unsaturated in composition. This may reflect underlying biosynthetic processes or the regulation of membrane biophysical properties subject to lipid substitution. Finally, several diglycosylceramide lipids increased up to ten-fold under P stress. These represent novel substitute lipids and potential biomarkers for the study of P limitation in situ, contributing to growing evidence highlighting the importance of sphingolipids in phycology. These findings contribute deeply to our understanding of P-lipid substitution, a powerful and widespread adaptation to P limitation in the oligotrophic ocean.
Importance

Unicellular organisms substitute phosphorus (P) containing membrane lipids with non-P substitutes when P is scarce, allowing greater growth of populations. Previous research with the model diatom species *Thalassiosira pseudonana* grouped lipids by polar headgroups in their chemical structures. The significance of the research herein is in the description of the individual lipids within the headgroups during P-lipid substitution. This revealed the relationship between lipid headgroups and hints at the underlying biochemical processes. Secondly, P-lipid substitution was contextualised by measurements of total cellular P. This depicted the place of P-lipid substitution in relation to the broader response to P-stress and yields insight into the implications of substitution in the marine environment. Finally, lipids previously unknown in this system were identified. This revealed a new type of non-P substitute lipid, potentially useful as a biomarker to investigate P-limitation in the ocean.

Keywords: *Thalassiosira pseudonana*, Phospholipid, Sphingolipid, Diatom, Lipidomics, Phosphorus, Stress, Limitation, Substitution, Biomarker.
Introduction

Diatoms are a diverse group of eukaryotic microalgae responsible for up to 25% of global and 50% of oceanic annual primary production (1). Lipids, hydrophobic or amphipathic biological molecules (2, 3), comprise 25 - 45% of the total dry weight of diatoms (4) representing a major pool of organic carbon. In the equatorial Pacific Ocean, for example, lipids account for 23% of the organic, total planktonic carbon (5).

Lipid biosynthesis involves a network of reactions that, through the exchange of polar headgroups and fatty acids, generate a rich variety of lipids (3, 6). Remodelling of a cell’s lipidome (the entirety of its cellular lipids) in response to environmental conditions is common in unicellular organisms (7, 8). One such remodelling mechanism utilizes the substitution of membrane glycerophospholipids with non-phosphorus glycerolipid counterparts when an organism is subjected to phosphorus (P) stress or starvation (7–10). This response allows a phytoplankter to reduce its P demands in P-limited environments (9), such as the Sargasso Sea or the eastern Mediterranean Sea. These oligotrophic regions contain very low bioavailable phosphate (PO$_4^{3-}$) concentrations (typically <10 nmol L$^{-1}$)(9, 11, 12). The reduction in total P demand confers a considerable advantage to growth by allowing for the prioritisation of non-substitutable functions, such as nucleic acid synthesis, over phospholipid biosynthesis (9, 13).

The marine diatom *Thalassiosira pseudonana* has been used as a model organism to study the effects of P starvation on lipid remodelling (8, 9). Under phosphorus replete growth conditions (P+) *T. pseudonana* synthesizes glycerophosphatidylcholine (PC), glycerophosphatidylglycerol (PG) and glycerophosphatidylethanolamine (PE) (8, 9). In contrast, when grown under P- conditions it synthesises the nitrogen containing betaine lipid diacylglycerylcarboxyhydroxymethylcholine (DGCC), which is normally undetectable under P+.
The increase in DGCC is concomitant with a decrease in PC and it is thought that the two physicochemically similar zwitterionic lipids can substitute for each other without loss of membrane function (8, 9, 14). Betaine lipids, including DGCC, are highly abundant in the marine environment (9, 15–18). In addition to the shift between PC and DGCC lipids in P-starved *T. pseudonana*, PG lipids may be exchanged for sulphur-containing sulfoquinovosyldiacylglycerol (SQDG) (8, 9). As such, ratios of the substitute lipid pairs have been considered as biomarkers of P stress in both marine and freshwater environments (9, 19).

The biosynthetic pathways underlying these processes are well defined for eukaryotic microalgae in general (20–23) and are coupled with the published genome of *T. pseudonana* (24). The biosynthetic pathway leading to DGCC, is conspicuously unknown, beyond the observed incorporation of radiolabelled methionine (25).

PG, and the glyceroglycolipids SQDG, monogalactosyldiacylglycerol (MGDG) and digalactosyldiacylglycerol (DGDG) are enriched in the plastid thylakoid membranes (26). Furthermore, these glyceroglycolipids are biosynthesised within the chloroplast via the intermediate diacylglycerol (DAG). The biosynthesis of cellular membrane lipids including PC and PE is, in contrast, conducted within the endoplasmic reticulum but also proceeds via a DAG intermediate (20–23). As such, the observed total cellular DAG composition may yield insight into lipid metabolism.

Lipid substitution kinetics in *T. pseudonana* are rapid, resulting in the exchange of the majority of cellular glycerophospholipids with DGCC and SQDG within 48 h of the initiation of P stress (8). P-starved *T. pseudonana* responds to the resupply of P faster still, restoring the predominance of glycerophospholipids over a 12 - 24 h period (8). P-lipid substitution dynamics have also been examined in other phytoplankton, such as the pennate marine diatom...
Phaeodactylum triconutum, resulting primarily in PG to SQDG and PC to diacylglyceryl hydroxymethyltrimethyl-β-alanine (DGTA) substitution, reflecting a contrasting system (27). In addition, a comprehensive recent study in Emiliania huxleyi illustrates increased DGCC:PC and SQDG:PG ratios in addition to ultrastructural modifications resulting from P stress (28).

P-lipid substitution in T. pseudonana has, therefore, been well characterised in terms of the total lipid within each of the major polar headgroup classes. However, important unknowns remain:

The dynamics of the individual lipid chemotypes, differentiated by the fatty acids they bear, subject to P stress remain unknown. Revealing these dynamics has the potential to yield insight into mechanisms of P-lipid substitution such as the synthesis of DGCC. Secondly, the effects of P-lipid substitution upon particulate organic P (POP) content per cell have not been reported. Consequently, the relationship between the responses associated with lipidic and extra-lipidic POP remains unknown. Moreover, previous work has not provided a comprehensive discussion of the relative contributions of two potential underlying mechanisms for P-lipid substitution, i.e. active break down of glycerophospholipids and replacement with non-P alternatives vs a simple switch in biosynthesis to the production of non-P lipids (29). Finally, the behaviour of minor lipid species, outside of the predominant headgroup classes, has yet to be studied. The cellular response to P stress is complex and powerful (30). It is expected therefore, that lipids other than those most common and abundant groups studied to date, are also affected. These minor species may display novel substitute or biomarker behaviour.

Presented herein, the findings of a mass spectrometry based lipidomics study, using both targeted and untargeted methodologies to characterise the lipidic response to P stress in cultures of T. pseudonana. These methods, coupled with culture growth monitoring, dissolved, and
particulate nutrient concentrations, yielded novel insight into the kinetics of P-lipid substitution, in relation to the broader cell response to P stress. In addition, the data on individual lipid species indicated the declining PC lipid pool became relatively enriched in unsaturated lipid species. Finally, untargeted lipidomic screening revealed diglycosylceramide sphingolipids that displayed lipid substitute/biomarker behaviour in the P stressed cultures. These findings offer a deeper understanding of the dynamics of P-lipid substitution in marine phytoplankton, a fundamental mechanism for growth in P scarce seas, and its cellular implications.
Results

Phospholipid Substitution and Particulate Organic Phosphorus Dynamics

Figure 1

Statistical significance of comparisons is indicated in the following text as *p<0.05 **p<0.005 by two-tailed, paired equal-variance T-test.

During exponential growth of the cultures (Figure 1A) between 0 and 72 h, the P+ cultures grew at a rate of 0.029 ± 0.0013 h⁻¹ (23.76 ± 1.08 h doubling time). Exponential growth during the same period in the P- cultures was marginally slower at 0.026 ± 0.0016* h⁻¹ (27.09 ± 1.78 h doubling time). Both the P+ and P- cultures appeared to be transitioning into stationary phase at 96 h with maximum populations of 1.08 x 10⁶ ± 4.11 x 10⁴ cells mL⁻¹ and 1.14 x 10⁶ ± 7.35 x 10⁴ cells mL⁻¹ respectively. Culture viability was high throughout at >90 %, in both P+ and P- cultures, with the exception of the P- cultures at 48 h (87.95 ± 1.78 %) (Supplementary Figure 1E), the cultures were therefore considered to be healthy and not subject to stresses other than P up to 72 h.

Dissolved phosphate ([PO₄³⁻], Supplementary Figure 1D) in the phosphorus replete (P+) control cultures was in excess (>10 µmol L⁻¹) throughout the 96 h course of the experiment. [PO₄³⁻] in the phosphorus stressed (P-) cultures at 0 h was 0.80 ± 0.00 µmol L⁻¹, dropping to 0.20 ± 0.00 µmol L⁻¹ after 6 h. This indicates a minor carryover of P from the P replete seed cultures to the P- culture treatment, which was depleted around 6-12 h thus subjecting the P- cultures to P
stress. Dissolved nitrate ([NO$_3^-$]) and silicate (orthosilicate ions [SiO$_4^{4-}$]) concentrations remained in excess throughout, in both P+ and P- cultures (Supplementary Figure 1A, B), with the exception of the P+ cultures after 72 h in which [SiO$_4^{4-}$] was depleted. The cultures were, therefore, not subject to confounding macronutrient stress, with the possible exception of the P+ 96 h samples. Taken together, the growth and nutrient data suggest the 96 h P+ samples were entering an Si limited stationary growth phase. These data will therefore, not be considered as a basis for drawing conclusions about P stress hereafter, but the data are included for context.

Particulate organic phosphorus (POP) quantity per cell (Figure 1B) fell with the progression of P stress through time, in the P- cultures. Initial P- POP was 1.94 ± 0.26 fmol cell$^{-1}$ and dropped consistently over 96h to a minimum of 0.25 ± 0.028 fmol cell$^{-1}$ at 96 h, equivalent to a 0.11 ± 0.03** fold reduction relative to the concurrent P+ control samples. The determined POP quantities per cell for the P+ cultures of between 1.79 ± 0.07 and 2.84 ± 0.22 fmol cell$^{-1}$ are approximately in line with previous observations for T. pseudonana of 2.8 – 6.2 fmol cell$^{-1}$ (31).

Total phospholipid (Figure 1C) represents the sum of the predominant phospholipid classes: glycerophosphatidylcholine (PC), glycerophosphatidylglycerol (PG), and glycerolphosphatidylethanolamine (PE). The individual dynamics of these lipid classes are shown in Supplementary Figure 2. Total P-lipid showed an initial increase in both P+ and P- cultures between 0 and 6 h, from 0.73 ± 0.08 to 1.25 ± 0.05 fmol cell$^{-1}$ in the case of the P- cultures and declined rapidly thereafter to a minimum of 0.17 ± 0.02 fmol cell$^{-1}$ at 96 h, equivalent to a 0.08 ± 0.01 fold reduction relative to the P+ control cultures.

The betaine lipid Diacylglycerylcarboxyhydroxymethylcholine (DGCC, Figure 1D), was absent from the P+ cultures throughout and the P- cultures before 12 h. Concomitant with the
decline in P-Lipid, DGCC quantities increased through time in the P- cultures following 12 h to reach a maximum of 0.85 ± 0.075 fmol cell$^{-1}$ at 48 h and 0.67 ± 0.036 fmol cell$^{-1}$ at 96 h.

The sulfolipid sulfoquinovosyl-diacylglycerol (SQDG, Supplementary Figure 2E), was 1.31 ± 0.14$^*$ fold greater in quantity in the P- cultures than the P+ control at 6 h. SQDG quantities were then statistically similar in P+ and P- between 12 and 72 h, followed by a 0.71 ± 0.045** fold reduction in the P- cultures, relative to P+ at 96 h. SQDG therefore, did not vary significantly in a consistent manner in response to P stress and did not appear to act as a substitute lipid for PG in this case, as found in other systems (9).

Diacylglycerol (DAG, Supplementary Figure 2D) quantity per cell did not vary significantly between P+ and P- cultures, with the exception of a 0.58 ± 0.10$^*$ fold reduction in the P- cultures at 96 h. To our knowledge DAG has not been previously characterised in P stressed T. pseudonana cultures and did not vary significantly and consistently in response to P stress.

Neutral storage lipids were not quantified in this study which was focused on polar and phosphorus containing membrane lipids. However, for context, it is known that diatoms biosynthesise large reserves of (non-P containing) triacylglycerol (TAG) storage lipids subject to P stress (32). This leads to an overall increase in total glycerolipids in the similarly sized pennate diatom P. tricornutum, subject to P stress (27).

**P-Lipids vs. Non-Lipidic Particulate Organic Phosphorus**

The difference between total POP and total P-lipid (Figure 1E) was used to investigate the dynamics of the non-lipid associated (Non-Lipidic) POP within the overarching P stress scenario.
In the P- cultures, the non-lipidic POP declined rapidly from 1.21 ± 0.27 to 0.083 ± 0.17 fmol cell\(^{-1}\) over the course of the first 12 hours, remaining approximately constant thereafter. Non-lipid POP in the P+ cultures, in comparison, remained statistically constant during the same time interval, decreasing over the course of 72 h from 1.19 ± 0.38 to 0.19 ± 0.12 fmol cell\(^{-1}\).

Degradation/Interconversion vs. Substitution by De novo Synthesis

The rates of change of P-lipid per cell and cell concentration were calculated (Figure 1F) to enable the distinction between a per cell ‘dilution’ whereby the original P-lipid was divided up amongst successive generations of progeny cells versus the chemical degradation and breakdown of the original P-lipid. The observations were made relative to the 12 h samples due to the initiation of P-lipid substitution at that time as evidenced by the depletion of dissolved P (Supplementary Figure 1D) and the commencement of DGCC synthesis (Figure 1D). The loss rate of P-lipid per cell of -0.031 ± 0.0015 h\(^{-1}\) was 32.29 ± 10.91**% faster than the dilution rate by culture growth of -0.023 ± 0.0015 h\(^{-1}\) (Figure 1F). The additional loss at greater than the rate of cell dilution is indicative of degradative breakdown.

In addition, the difference in total phospholipid per unit volume culture, between time (t) and 12 h in the P- cultures (\(\Delta PL_t-12h\)) was also used to assess the total quantity of lipid per culture, regardless of per cell quantities (Supplementary Figure 2F). At 48 and 72 h in the P- cultures, \(\Delta PL_t-12h\) was -42.27 ± 18.19* pmol mL\(^{-1}\) and -80.47 ± 20.03** pmol mL\(^{-1}\) respectively, indicating a net degradation of the total phospholipids at these times. The P liberated from P-lipids between 12 and 72 h was 7.66 x 10\(^7\) ± 1.95 x 10\(^7\) atoms per daughter cell grown during that time, equivalent to 1.06 ± 0.55 diploid genomes per cell (8, 33). Therefore, the P released from P-lipid
breakdown appeared sufficient to cover DNA biosynthesis requirements for growth during this time interval.

The maximum degradative decrease in ΔPL₄₋₁₂h at 72 h (Supplementary Figure 2F) corresponded to a loss of 34.10 ± 8.75 % of the maximum total phospholipid observed at 12 h in the P- cultures. This value is consistent with the P-lipid degradative loss rate calculated previously (32.29 ± 10.91** % faster than the dilution rate by culture growth) over the same time interval. In comparison, the change in total DGCC (DGCC₄₋₁₂h) increased rapidly through time reaching +464.90 ± 50.05 pmol mL⁻¹ at 72 h. Taken together, the molar degradative decrease in PL₄₋₁₂h was equivalent to 17.31 ± 4.69 % of the increase in DGCC.
Variation in individual lipid species quantity, between P+ and P- cultures at 72 h is shown in Figure 2. All PC species (Figure 2A) showed a dramatic decrease in quantity per cell in the P- cultures, by comparison to the P+ control cultures. Some variation in the magnitude of this decrease was evident from the data: PC(18:4/16:0) and PC(20:5/18:4) showed the largest fold decreases of 0.018 ± 0.0035** and 0.023 ± 0.037** fold respectively. PC(16:1/16:0), PC(20:5/16:1), PC(20:5/16:0) and PC(38:6) showed highly significant, intermediate fold decreases between 0.053 ± 0.011 and 0.15 ± 0.043. In contrast, highly unsaturated PC(22:6/20:5) and PC(20:5/20:5) species displayed the smallest fold decreases of 0.28 ± 0.054** and 0.30 ± 0.052** fold respectively.

Betaine lipid DGCC species (Figure 2B) occurred only in the P- cultures and were below detection in the P+ control cultures. At 72 h, the five most abundant DGCC species in the P- cultures were DGCC(20:5/16:0), DGCC(22:6/20:5), DGCC(20:5/20:5), DGCC(38:6) and DGCC(20:5/16:1), with quantities of 84.28 ± 14.14, 60.08 ± 9.39, 56.56 ± 11.87, 50.56 ± 7.40 and 33.05 ± 6.16 amol cell⁻¹ respectively.

Glycerophospholipid PE species (Figure 2C) decreased in the P- cultures relative to the P+ control cultures. The predominate PE lipid species PE(22:6/20:5), PE(20:5/20:5) and PE(20:5/16:1) decreased by 0.19 ± 0.065**, 0.18 ± 0.060* and 0.07 ± 0.028** fold respectively under P stress.
The glycerophospholipid PG (Figure 2D) behaved in line with PC and generally showed a decrease in quantity per cell for its molecular species in the P- cultures, relative to the P+ control cultures. The predominant chemotypes, PG(16:1/16:0)**; PG(20:5/16:0)*; PG(20:5/16:1)**; PG(16:1/14:0)**; and PG(32:2)** all exhibited fold decreases of between 0.062 ± 0.027 and 0.11 ± 0.048 subject to P stress.

Bivariate analyses (Supplementary Figure 3) were used to compare the relative abundance of individual lipid species within pairs of lipid classes, in order to provide an indication of similarity in their fatty acid distributions. Overall, when considering the distributional similarity between all the measured lipid classes in both treatments, at all time points between 12 and 72 h, two primary clusters were observed: The first cluster was comprised of PC, its substitute DGCC, and PE and bore similar fatty acid distributions with an average pairwise correlation coefficient (r) of 0.68 ± 0.25. A second cluster, of PG, DAG, MGDG and DGDG fatty acid distributions was highly cross correlated with r = 0.87 ± 0.10. In addition, these two clusters were highly distinct from each other, with r = 0.074 ± 0.18. Interestingly, DGCC was moderately correlated with DAG at 12 h, with r = 0.53 ± 0.023 and weak to no relationship thereafter. DGCC abundance at 12 h was, however, negligible compared to 24 h and later.

The absolute quantities of degradative loss of PC individual lipid species were weakly correlated with DGCC gains (Supplementary Figure 4D, E and F) for time intervals of t48-t24 h and t72-48 h in the P- cultures, with coefficients of determination between $R^2 = 0.3^{**}$ and 0.17* respectively. The linear regression coefficients were consistent with 18 and 20% respectively of the DGCC synthesised at these time intervals originating as recycled diglyceride moieties from degraded PC. This estimate is further consistent with the cumulative degradation of PC between 12 and 72 h, presented previously, of 17.31 ± 4.69 % of the increase in DGCC at the total class level (Supplementary Figure 2F).
Fatty Acid Level Variability during P-replete Growth

In order to study the temporal dynamics of P-lipids in the context of P stress at the individual lipid species level, the top 5 most abundant PC lipids, as the predominant components of the total P-lipid pool, were analysed in isolation. Firstly, the underlying dynamics of the P+ control cultures were considered (Figure 3A). The top five PC molecular species, as ranked by their maximum quantity per cell, appeared to conform to one of two behaviours. The highly unsaturated PC(20:5/20:5) and PC(22:6/20:5) chemotypes displayed a sharp increase between 0 and 12-24 h with the initiation of exponential growth, followed by an equally sharp decline in abundance to 72 h. PC(22:6/20:5), for example, rose from 18.11 ± 4.39 at 0 h to 168.57 ± 9.83** at 24 h, before dropping back to 39.67 ± 7.28** amol cell⁻¹ by 72 h. In contrast, moderately unsaturated species PC(18:4/16:0), PC(20:5/18:1), and PC(20:5/18:4) displayed a consistently increasing trend with the progression of time in the P+ control cultures. PC(20:5/18:4), for example, increased from 36.21 ± 7.44 at 0 h to 159.26 ± 39.81* amol cell⁻¹ at 96 h.

To decouple the P stress driven dynamics from the growth driven dynamics, the change in PC lipid quantity through time, relative to the initiation of P stress in the P⁻ cultures at 12 hours was determined (Figure 3B). Overall, the top 5 PC lipids all exhibited a continuous decreasing trend in quantity between 12 and 96 h subject to P stress. The magnitude of change was variable between different lipid species, with PC(20:5/18:4) and...
PC(20:5/18:1) decreasing by 0.52 ± 0.081** and 0.16 ± 0.025** fold respectively. The response of PC(22:6/20:5) was an exception, showing an initial 1.46 ± 0.11 fold increase between 12 and 24 h, followed by a large 0.33 ± 0.073** fold decrease (relative to 12 h). The least unsaturated of the major PC lipids, PC(18:4/16:0) showed the most rapid initial decrease between 12 and 24 h dropping by 0.64 ± 0.070 fold.

The variability in the response of these PC lipids to P stress was reflected in the changing average unsaturation of the PC lipid pool through time (Figure 3C). Average PC unsaturation (the total number of fatty acid double bonds) in both P+ control and P- cultures displayed a sharp increase in average PC lipid unsaturation between 0 and 12 h, from 5.73 ± 0.098 to 7.53 ± 0.24 in the P+ case. After 12 h, the results diverged between the treatments, concomitant with the initiation of P stress. The P+ cultures then displayed a decrease in average PC unsaturation to 5.63 ± 0.077 at 72 h, remaining constant thereafter. The P- cultures, in contrast, continued increasing to a maximum of 7.95 ± 0.14 at 24 h, followed by a decline until 96 h to 6.37 ± 0.25, but remained more highly unsaturated than the P+ control cultures throughout.

These observations co-occur with the P- cultures experiencing decreasing per cell quantities of P-lipids and a minor degree of degradative breakdown as described herein (Figure 1C). Therefore, the initial shift to higher unsaturation in the P- cultures indicates that P stress causes either a preferential degradation of less saturated PC lipids or the continued synthesis during early P stress of low quantities of highly unsaturated PC lipids between 12 and 24 h (both mechanisms appear supported by Figure 3B). At later time points, this ceases and the average PC unsaturation begins to decrease and trend toward (but not achieve) convergence with the P+ control values. Overall, a more highly unsaturated PC lipid pool occurs subject to P stress (Figure 3C).
Untargeted Lipidomic Screening

Figure 4

Lipid species were ranked according to their normalised abundance, in descending order from the most strongly increased under P stress (Figure 4B). The betaine lipid diacylglycerolcarboxyhydroxymethylcholine (DGCC) dominated the positive ion results comprising 14 of the top 15 ions. DGCC species containing eicosapentaenoic (20:5), docosahexaenoic (22:6) and palmitic (16:0) fatty acids, displayed the greatest differential increases and were hence the highest ranked.

Most remarkably, a 1014.7112 Da species, corresponding to a diglycosylceramide with a dihydroxy(18:3) long chain base and a 24:0 fatty amide ((Gly)$_2$Cer(d18:3/24:0)) with a formic acid adduct, displayed an absence-presence response to P stress. The chemical structure, and its determination by MS2 fragmentation in both negative and positive ion mode, is displayed in Supplementary Figure 5.

Figure 5

Several lipids closely related to (Gly)$_2$Cer(d18:3/24:0) were subsequently observed using more sensitive semi targeted analyses in positive ion mode (Figure 5). Of those, (Gly)$_2$Cer(d18:3/24:0) itself responded the strongest to P stress with a 10.43 ± 3.12** fold increase between the P- and P+ cultures. Each of the glycosphingolipids displayed a
(Gly)$_2$Cer(d18:3/24:0), (Gly)$_2$Cer(d18:2/24:0) and (Gly)$_2$Cer(d18:1/24:0) increased by 6.67 ± 1.81**, 4.89 ± 1.76** and 2.03 ± 0.36** fold respectively, between the P- and P+ cultures. The ceramide equivalent lipid Cer(d18:3/24:0), without the diglycosyl headgroup, was also detected. In contrast, its differential abundance between P- and P+ cultures showed no significant change.
Phospholipid Substitution and Particulate Organic Phosphorus Dynamics

We find that the dynamics of P-lipid substitution in *T. pseudonana* follow primarily a cessation of P-lipid net biosynthesis with a minor degree of P-lipid net breakdown (Figure 1). The P liberated by degradative breakdown of P-lipids was enough to synthesise approximately one diploid genome per cell during the same time period of P-stressed growth, one fourth of previous estimates (8). The majority of the original cellular P-lipids were retained and diluted in per cell terms with the progression of culture growth by cell division, consistent with the observed decrease in per cell POP. No appreciable growth limitation occurred on these timescales, reflecting the great capacity of this organism, not least with respect to its lipidomic flexibility, to adapt to low P environments (30).

Non-lipidic P per cell was observed to decrease to near zero after 12 h and warrants discussion, given the expected obligate cellular P requirements for essential genomic and other biomolecules. The minima in non-lipidic P of 0.076 ± 0.058 fmol cell\(^{-1}\) at 24 h is just barely sufficient to account for the estimated P associated with DNA per cell (63.74 ± 48.64% estimated from (33)), when the upper standard deviation limit is included. Therefore, it is possible that the measured total POP quantities represent a slight underestimate. This is supported by the measured P+ POP per cell quantities, which are shown in the results section to be consistent with the lower limits of literature values. Nonetheless, the relative shift from the initial state toward a minimum value after 12 h, remains a robust observation appropriate for further discussion.

Non-lipidic POP declined toward its minimum by 12 h, prior to the initiation of the P-lipid response. The temporal delay in the lipid remodelling response to P stress may be attributable to the prior utilisation of P from other intracellular pools. Polyphosphates (polyP)
could be one such pool (34), however these observations conflict with recent environmental observations which suggest polyP enrichment of phytoplankton in P-limited waters (35). To our knowledge, polyP has not been quantified in *T. pseudonana* to date, so this contradiction cannot be reconciled at this point.

The majority of the original lipid bound P remained as such and did not appear to be made available for other cellular processes such as nucleic acid synthesis (8, 27). This does not diminish the ability of the organism to grow subject to reduced P availability by utilising non-phosphorus containing lipids rather than phospholipids. The distinction has subtle conceivable ecological implications. Heterotrophic bacteria that consume lipids efficiently (36), for example, may benefit from access to lipidic P content in oligotrophic waters. Additionally, assuming that the use of non-phosphorus containing lipids rather than phospholipids has little physiological influence on cell growth, the phospholipid content of P replete cells arguably represents an enigma if this P pool is not strongly re-purposed on development of P-stress.

The stable abundance of SQDG throughout could be interpreted as an adaptation to economize on P requirements in the oceanic environment in which P is regularly scarce. The substitute lipid pair, PG and SQDG, share at least a partial functional similarity/redundancy in their role in the thylakoid membranes (10, 26). However, the efficiency of photosystem II is demonstrably compromised in SQDG deficient algal mutants (37), suggesting SQDG has an essential functional role and is not only a non-P substitute for PG.

Overall, this deeper understanding of a powerful phytoplankton adaptation to P-stress gives greater context to the resulting dramatic shifts upon elemental ratios and phytoplankton growth dynamics, with subsequent implications to both trophic transfer and export flux.
Fatty Acid Level Response to Phosphorus Stress

PC individual lipid quantity per cell was highly dynamic in the P+ control cultures with the progression of the time (Figure 3A). The most highly unsaturated PC lipids increased markedly in early exponential growth before declining back to initial levels later as the cultures progressed toward stationary growth phase. More saturated major PC lipids displayed a progressively increasing quantity per cell through time. These observations are in agreement with previous reports on the relationship between unsaturation of eukaryotic phytoplankton glycerolipids and growth phase (38). The change in lipid unsaturation state arises from the partitioning between polar and neutral storage lipid biosynthesis, which proceed via distinct elongation and desaturation pathways (38). When growing rapidly, polyunsaturated fatty acids characteristic of the structural polar lipids are biosynthesised. However, during the transition to stationary growth, more saturated fatty acids are synthesised for incorporation into TAG storage lipids. This is reflected in part in the polar glycerolipids due to a degree of crosstalk (38).

The discussed variation in membrane glycerolipid composition should have subsequent functional implications upon membrane biophysics and structure, unless homeostatically compensated in some other manner (39). The complexity and degree of the temporal changes in the individual lipid species composition, within the P+ control cultures, highlight the lipidomic plasticity of T. pseudonana and its highly dynamic nature, both often poorly accounted for in biomarker studies. This behaviour contributes to a lack of diagnostic power for chemotaxonomic distinction of phytoplankton that has been reported elsewhere with respect to fatty acids (40) and intact polar lipids (18).

In the P- cultures, we observe a preferential degradation of more saturated PC lipid species and/or a continuing synthesis of highly unsaturated PC lipids, leading to a more
highly unsaturated, less abundant PC lipid pool (comprising the majority of the P-lipids) under P stress (Figure 3B, C). It could be speculated that this increase in unsaturation represents a homeostatic control on membrane fluidity (as in (39)) in light of the increasing substitution of P-lipids by DGCC, which may result in differing biophysical characteristics.

DGCC, which increased in cellular abundance with the progression of P stress, was correlated in chemical composition with PC (Figure 2 and Supplementary Figure 3). This similarity in composition is consistent with the role of DGCC as a substitute lipid for PC in T. pseudonana (8, 9), and the observation provides further evidence for the substitutive link between the two.

The molar quantity of P-lipid breakdown accounted for a minor proportion of the biosynthesised DGCC at the total class level. In addition, the degradative loss of individual PC lipid species was weakly correlated with the synthesis of their DGCC counterparts (Supplementary Figure 4). These data are consistent with, but not definitive evidence of, the recycling of diglyceride moieties from phospholipid breakdown for incorporation into newly biosynthesised DGCC. This observation must be further confirmed by an isotopic labelling experiment to unambiguously trace the transformation.

DAG is a known direct precursor in eukaryotes to glycerophospholipids (PC, PE)(20). Under P- conditions, an uncoupling of the DAG composition with the glycerophospholipids could be expected if they are not being synthesised. However, DAG composition did not vary between P+ and P- conditions as previously discussed. Therefore, we expect the composition of lipid species in the DAG to reflect that of PC, the most abundant of its biosynthetic products, under P+ conditions but that was not the case. Furthermore, no consistent correlation was observed between DGCC and DAG. As the precursor to PC biosynthesis (20) and a substructure in common to both PC and DGCC, DAG is a potential intermediate in the
synthesis of DGCC. This could occur via de novo synthesis or represent a recycling of any diglyceride from PC breakdown via phospholipase C. Again, our observations do not support these hypotheses. PC is synthesised from DAG under P+ conditions, yet we cannot observe this via the correlation of its individual lipid species. Based on these results, a role of DAG in the biosynthesis of DGCC cannot be ruled out. Unfortunately, comparison of the various polar lipid distributions did not yield significant insight into the as yet unknown biosynthetic pathway leading to DGCC. To our knowledge, nothing appears known other than the reported incorporation of 14C labelled methionine into its head group in labelling experiments (25, 41). This remains a very conspicuous gap in our understanding of a lipid class that forms a major component of the planktonic lipid pool (42).

Thus, it appears that the observed DAG is utilised primarily by monogalactosyldiacylglycerol synthase as a biosynthetic precursor to MGDG and DGDG (21, 24). These observations lead to the formation of the hypothesis that there are two (or more) separate pools of DAG. Firstly, the larger and/or slower turned over DAG pool indicative of MGDG/DGDG synthesis, observed by characterisation of the total lipid extract and localised within the chloroplast. Secondly, the smaller and/or more rapidly turned over DAG pool indicative of PC/PE synthesis that is conspicuously not observed. Further investigation of DAG lipid dynamics within isolated subcellular locales may yield further insight into the link between DAG and the biosynthesis of PC and DGCC.

Untargeted Lipidomic Screening

Untargeted lipidomic screening revealed (Gly)$_2$Cer(d18:4/24:0) which was approximately 10 fold more abundant in the P- than the P+ cultures. This discovery led to the identification of a series of related molecules varying subtly in the degrees of unsaturation of the long chain
base or the fatty acid amide (Figure 5). All of the diglycosylceramide species increased under P stress, varying in magnitude of change between approximately 2 and 10 fold. A Cer(d18:3/24:0) species was also identified and demonstrated contrasting behaviour. This putative precursor did not vary in cellular abundance subject to P stress. Chemically identical and related lipids have been previously reported in another marine diatom, Skeletonema costatum (43) and genes encoding sphingolipid biochemistry are also annotated in the genome of T. pseudonana (24). This lends further credence to our identification and highlights these molecules as an area of interest for future research into diatom lipid biochemistry. Glycosylceramides have been ascribed to several physiological functions including membrane stability, membrane permeability and pathogenesis (44). In this context, it appears that they may be acting as non-phosphorus substitute lipids, akin to DGCC (8–10), for another phosphorus containing lipid and this drives their differential increase under P stress.

Chemically closely related glycosphingolipids have been identified in virally infected Emiliania huxleyi and applied as biomarkers for viral infection in the marine environment (45–48). In the same manner, we propose the diglycosylceramides that increase in cellular abundance subject to P stress, as candidate biomarkers for the P stress of marine diatoms in the environment. As such, the abundance of these potentially diagnostic biomarkers could be quantified in diatoms collected from the environment of interest. This data could yield additional insight into the level of P stress experienced by the phytoplankton directly, which would be complimentary information to the more routine measurement of dissolved or particulate P concentrations in the medium. Targeted quantification of these lipids would be readily achievable using commercially available lactosyl-ceramide standards as in (43). Such a lipid biomarker could be useful to follow P related biogeochemical processes in the environment, pending further validation. More broadly, the discovery of this sphingolipid
behaviour, taken together with the *E. huxleyi* host/virus system strongly supports phytoplankton sphingolipid biochemistry as an important avenue for future research.
Conclusions

We have presented herein the comprehensive lipid analysis of phospholipid substitution induced by P stress in the model marine diatom *T. pseudonana* at the level of individual lipid species, those varying in fatty acid composition, and in tandem with quantification of cellular particulate organic phosphorus (POP). These data indicate depletion of a non-lipidic POP fraction, prior to the lipid response. This was followed immediately thereafter by de-novo biosynthesis of the substitute lipid DGCC. The majority of the original P-lipid remained intact and was diluted amongst progeny cells but P equivalent to approximately one diploid genome per daughter cell was liberated by degradative breakdown of P-lipids. The distinction between P associated ultimately with lipids as opposed to genomic material may have implications upon subsequent trophic transfer and export flux.

At the individual lipid level, all major P-lipids reduced in cellular quantity but in a non-uniform manner. P stress resulted in an increased average degree of unsaturation within the declining PC lipid pool, which represented the predominant P-lipids. These observations indicate a preferential degradation of more saturated PCs and/or a continuing synthesis of highly unsaturated PC lipids, perhaps in order to regulate membrane fluidity in the context of the major change in membrane composition through substitution. Further, the data were tentatively consistent with the recycling of diglyceride substructures from P-lipid degradation into newly synthesised DGCC.

DAG fatty acyl composition primarily reflected that of the chloroplast associated glyceroglycolipids and did not yield insight into the biosynthetic pathway of the substitute lipid DGCC. This pathway remains a significant gap in our understanding of a biochemical highly abundant in the marine environment.
Finally, untargeted lipidomic screening revealed a group of diglycosylceramide lipids which increased subject to P stress up to 10 fold and constitute non-phosphorus substitute lipids and candidate biomarkers for P stress. Taken together, these findings contribute a new level of understanding of P-lipid substitution in marine phytoplankton, a powerful and widespread adaptation to low P environments with important consequences upon macronutrient biogeochemistry and oligotrophic primary production.
Experimental Procedures

Culturing

Axenic *T. pseudonana* (1085/12 also designated CCMP1335/3H) was obtained from the Culture Collection of Algae and Protozoa, Scottish Association for Marine Science, U.K. Culture manipulations were performed under sterile, laminar flow environment.

F/2+Si growth media (49), based on artificial seawater was prepared (50) from analytical or biological grade components (Fisher Scientific). Seed culture (175 mL) was grown to mid-log phase concentration of $1.13 \times 10^6$ cells mL$^{-1}$ over 4 days incubation at 18°C; 12:12h light/dark cycle; 123 µmol quanta m$^{-2}$ s$^{-1}$ illumination; 70 rpm gentle orbital agitation.

The seed culture was split (2 x 79.6 mL) and cells isolated from the media by filtration (Millipore Steritop, 0.22 µm pore size). Cells were washed on the filter with 50 mL of P+/P$^-$ media, depending on treatment, then resuspended and split to form 3 x 300 mL for each P+ and P-. These experimental cultures were incubated as above and sampled after 0; 6; 12; 24; 48; 72 and 96 hours for size distribution/cell count; viability; dissolved/particulate macronutrients and lipid extracts.

An aliquot (950 µL) of culture was mixed with freshly prepared paraformaldehyde solution (170 µL, 34% w/v, dH$_2$O) and stored at 4°C for <24 h before analysis. Cell size distributions were generated with a Beckmann Coulter Multisizer 3 Coulter Counter. A 70 µm aperture and 3% NaCl electrolyte were used and the samples diluted to ensure <10 % aperture coincidence concentration. The Coulter Counter was calibrated with 5.023 µm polystyrene latex standard beads prior to use (Beckmann Coulter via Meritics Ltd., Dunstable, U.K.). Size distributions were used to generate cell concentration values, between the limits of 3 and 9 µm particle diameter.
Experimental culture (50µL) was incubated with SYTOX-Green dye (Invitrogen Life Technologies, Paisley, U.K.) at a concentration of 0.5µM for 5 minutes in the dark. 18 µL of this solution was then imaged with a Cellometer Vision Duo (Nexcelcom Bioscience via. Peqlab, Sarisbury Green, U.K; X100-F101 Optics; SD100 Slides). All cells were counted manually under brightfield mode and stained, non-viable cells under fluorescence mode (470/535nm excitation/emission).

**Nutrient Quantification**

Experimental culture (10 mL) was syringe filtered over pre-combusted (450°C, 12h) GF/F filters. The filtrate was stored at -20 °C. Filters were dried at 60°C for 24 h and stored in a desiccator. Particulate phosphorus was determined following an oxidation procedure as described in reference (51), samples were centrifuged (1000 x G, 10 mins, 18 °C) before sampling to remove particulates. Nutrient samples were diluted: 1/60 (dissolved nutrients (filtrate)) and 3/20 (oxidised particulate nutrients) in milliQ dH2O and characterised by segmented flow autoanalysis on an AutoAnalyzer 3 (Seal Analytical, Fareham, U.K.) for phosphorus, nitrate/nitrite and silicon. POP quantities were corrected for carryover of inorganic phosphate from the growth media based upon the difference between the P+ and P- POP measured at 0 hours, normalised to the $[\text{PO}_4^{3-}]$ concentration in the P+ growth media for a given sample (as measured by the dissolved phosphate (DIP) assay):

$$((\text{PO}_{P+}^{t=0})-(\text{PO}_{P-}^{t=0})) \times ((\text{DIP}_{P+}^{t=0})/(\text{DIP}_{P-}^{t=0}))$$

**Lipid Extraction**
Lipid samples (20 mL) from the experimental cultures were isolated by syringe filtration as above. The filtrate was discarded and the filters stored at -78 °C until extraction. Total lipid extracts were prepared using a Bligh-Dyer extraction procedure (52) as modified by (53). Solvents were LC-MS grade (Fisher Scientific). Lipid internal standards were added at quantities adjusted per sample to maintain constant standard:cell equivalents ratios as follows: PC(12:0/12:0)(56.2 amol cell\(^{-1}\)); PG(12:0/12:0)(77.0 amol cell\(^{-1}\)); PE(12:0/12:0)(47.6 amol cell\(^{-1}\)); DAG(20:4/18:0)(99.7 amol cell\(^{-1}\)) and SQDG (mixed extract, predominantly 34:3)(1.27 fmol cell\(^{-1}\)). Phospholipid/DAG standards were acquired from Avanti Polar Lipids (Alabaster, U.S.A.), SQDG spinach leaf extract was provided by Lipid Products (Surrey, U.K.).

A variable volume of this phase was isolated per sample ensuring a constant quantity of lipid cell equivalents/standard quantity was infused into the mass spectrometer during analysis (hence accounting for ion suppression and enabling external standard quantification of DGCC). The optimal quantity for analysis under instrument/conditions below was found to be 0.8 x 10\(^6\) cells (+standards as described). This fraction was dried under N\(_2\) and stored at -20°C until analysis.

### Direct Infusion ESI-MS/MS Quantitative Analysis

Mass Spectrometric analysis was performed on a Waters Micromass Quattro Ultima triple quadrupole instrument. Dried samples were dissolved in 250µL 66% methanol, 30% dichloromethane, 4% ammonium acetate (300 mM in H\(_2\)O). The sample solution was directly infused into the instrument at 6µL min\(^{-1}\) and analysed by MS/MS analysis of each lipid class (53). In addition, a neutral loss scan of 35 Da, between 350-750 Da at a collision energy of 15 eV, was used for the detection of ammoniated DAG molecular ions.
Spectra were processed by despiking, baseline subtraction, isotopic correction and assignment by a visual basic macro (54). The spectrum of the SQDG standard in isolation was taken in triplicate and used to perform a subtraction for overlapping standard derived peaks (due to the natural extract nature of the SQDG standard) based on the dominant peak at 834 Da (SQDG34:3, not detectable in *T. pseudonana* under P+/P- conditions). DGCC extract (purified by preparative HPLC from *T. pseudonana* under P starvation) was provided by Benjamin A. S. Van Mooy (Woods Hole Oceanographic Institute, U.S.A.). External standard calibrations were generated from addition of DGCC extract (0.094; 0.19; 0.38; 0.75; 1.5 nmol) to P-replete grown (hence no intrinsic DGCC) *T. pseudonana* total lipid extracts prepared at optimal, constant cellular lipid/standard concentration as previously discussed. DGCC total counts were normalised to the chemically similar PC internal standard for quantification of experimental samples: Counts$_{DGCC}$/Counts$_{PC}$ = 18.594 x Quantity$_{DGCC}$. Highly linear over this range ($R^2 = 0.992$).

**Untargeted UPLC-ESI-AutoMS$^2$ Analysis**

UPLC-ESI-AutoMS$^2$ analysis was performed on a Dionex UltiMate 3000 UPLC system coupled to a Bruker maXis 3G quadrupole - time of flight (Q-ToF) mass spectrometer with an electrospray ionisation source. Lipid samples were dissolved in methanol (200 μL) prior to analysis. A 20 μL injection was taken by autosampler from vials in a cooled sample tray at 5°C. The sample was then chromatographically separated over 30 minutes with a Waters Acquity UPLC BEH C8, 1.7 μm particle, 2.1 x 100mm column. A constant flow rate of 0.3 mL min$^{-1}$ was used resulting in back pressures of between 260 and 460 bar.

Eluent A was water with 0.2% formic acid and 1% 1M ammonium acetate, eluent B was methanol with 0.2% formic acid and 1% 1M ammonium acetate. The column was heated
to 50°C and the eluent cooled to 21°C post-column, throughout the analysis. The following multi-step linear gradient was applied, with a constant flow rate of 0.3 mL minute\(^{-1}\): 35% eluent B at 0 minutes, increasing to 80% eluent B at 2 minutes, increasing to 95% eluent B at 12 minutes and holding for a further 18 minutes until the end of the run. Eluent B was decreased to 35% over 0.5 minutes post run and the column allowed to equilibrate for 4.5 minutes prior to the next run.

The mass spectrometer was calibrated by direct infusion of sodium formate solution prior to use (10 mM sodium hydroxide + 0.2% formic acid in 1:1 isopropanol/water). The observed mass accuracy was 0.4 ppm, determined from the standard deviation from the quadratic calibration curve for calibrant ions up to 1000 Da, in positive ion mode. The Q-ToF mass spectrometer yielded a mass resolving power of 21463.70, determined from full width at half maximum (FWHM) of the internal standard peak dilauroylphosphatidylcholine at 622.4470 Da [M+H]+ in positive ion mode. Full scan MS was acquired in positive and negative mode between 30 and 1500 Da.

During each run (positive and negative ion mode require separate analytical runs) ions above the noise level threshold were subjected to data dependant MS\(^2\) fragmentation. The threshold was set at 2000 and 1000 counts in positive and negative ion mode respectively. Ions with a mass of between 500 - 1500 and 300 - 1500 Da were subject to MS\(^2\) fragmentation in positive and negative ion mode respectively. The most abundant two precursor ions eluting during an MS scan were fragmented and after two MS\(^2\) spectra were acquired for a given ion, they were actively excluded from further MS\(^2\) for 1 minute. Fragmentation for MS\(^2\) was achieved by collision induced dissociation (CID) by impact with Argon gas, with stepped collision energies for precursor ions of increasing mass. Ions of 300-500, 500-800 and 800+ Da were fragmented with collision energies of 25, 40 and 50 eV respectively in positive ion mode and 25, 30 and 40 eV in negative ion mode.
Untargeted Data Processing

Bruker CompassXport was used to export the raw data prior to processing with the MZMine 2 software package (55). MZMine was used to generate extracted ion chromatograms and match these chromatograms between different samples. Integrated peak areas were normalised to the total number of cells extracted and adjusted for recovery of the internal standard (dilauroylglycerophosphatidylcholine). Peak assignments were based upon matching to an extensive, accurate mass, structure query language (SQL) lipid database generated in house. The database was populated by permutations of fatty acids (chain length/degree of unsaturation) and common glycerolipids/sphingolipids. The complete LIPID MAPS (version 20130306) structural database (56) and MaConDa mass spectrometry contaminants database (57) were also included.

The chemical formulae of database entries were then used to calculate accurate mass m/z values based upon a list of common molecular ion adducts in ESI-MS (58). Tentative assignments were made by matching precursor mass ions with the theoretical database to within a mass difference of < ±10 ppm. Database assignments were then confirmed by the identification of supporting MS2 fragments in each case. Fragments were assigned within a tolerance of < ±20 ppm unless otherwise stated.

The MS2 data was assigned using an in house visual basic macro, matching fragment ions to a database of common and diagnostic fragments and dynamically generated neutral losses. Matches were made based upon a tolerance of < 20 ppm (unless otherwise specified).
Acknowledgements

The authors would like to acknowledge John R. Gittins, Stephanie Tweed, Sophie Richier, Mark Stinchcombe and Victoria Goss for methodological assistance. We thank Benjamin van Mooy and Helen Fredricks (Woods Hole Oceanographic Institution, U.S.A.) for providing the DGCC standard. This work was funded by the University of Southampton - Vice Chancellors Scholarship Award. The purchase of the mass spectrometers was supported by the Wellcome Trust (Grant 057405).

Author Contributions

Jonathan E. Hunter designed the experiments, developed methods, carried out culturing, sample preparation, analysis, data interpretation and wrote the manuscript. The remaining co-authors assisted with experimental design, analysis, interpretation and drafting the manuscript.
References


31. Núñez-Milland DR, Baines SB, Vogt S, Twining BS. 2010. Quantification of


59

60
Figure Legends

Figure 1: Cell concentration growth curve (A); particulate organic phosphorus (POP) per cell (B); Total phospholipid (P-Lipid, C); total DGCC (D) and the quantity of non-lipidic POP per cell (E) in the phosphorus replete (P+) and phosphorus stressed (P-) cultures, under P+ and P- conditions, with the progression of time. Panel (F) depicts the loss rate of the total phospholipids (P-Lipid) in comparison with the cell dilution rate. Values are relative to 12 h, observed as the initiation of P stress and consequently DGCC biosynthesis and P-Lipid substitution/degradation in the P- cultures. Data are the mean of three biological replicates, with shaded error regions of one standard deviation.

Figure 2: Individual lipid molecular species quantity per cell within each of the lipid classes measured at 72 h in the P+ and P- cultures. All quantities are in amol cell$^{-1}$ with the exception of MGDG (G) and DGDG (H), for which relative, percentage abundances are shown. Molecular species contributing less than 5% of the total quantity in their lipid head group class in both treatments were excluded for brevity. The predominant fatty acyl combinations are shown where identified and consistent across the present lipids. Data are the mean of three biological replicates, with error bars of one standard deviation. Statistical significance is indicated in the figure and accompanying text, *p<0.05 **p<0.005 by two-tailed, paired equal-variance T-test.

Figure 3 – Top 5 most abundant individual PC lipid molecular species quantity per cell, through time, in the P+ control cultures only (A), change in the top 5 PC lipid molecular species, between time (t) and the initiation of P stress (12 h) in the P- cultures only (B) and the average unsaturation of the total PC lipid pool in P+ and P- treatments, with the
progression of time (C). Data are the mean of three biological replicates, with shaded error regions of one standard deviation.

Figure 4 – Untargeted lipidomic screening determined a total of 1161 different molecular ions in the 72 h P+ and P- cultures using positive and negative ion mass spectrometry. The normalised abundance in the P- cultures, indicating how strongly a given ion was associated with the P stressed cultures, is plotted against mass to charge ratio (m/z) and chromatographic retention time (R.T.) in panel (A). The top 15 ions associated with the P stressed cultures in positive and negative mode identified where possible by database matching and MS2 fragmentation analysis are shown in panel (B).

Figure 5 - Fold change, subject to P stress, in observed ions related to (Gly)$_2$Cer(d18:3/24:0), in positive ion mode. Long chain base and fatty acid amide fragments were observed in support of each of the following assignments in positive MS2. Data represent the mean of biological triplicate samples with error bars of one standard deviation. Assignments represent the primary fatty acyl configuration, as determined by the abundance of the fatty acyl fragments in the MS2 spectra.
(A) Lipids highly indicative of P- (in positive and negative ion mode):

<table>
<thead>
<tr>
<th>Lipid ID</th>
<th>Mode</th>
<th>Normalised Abundance</th>
</tr>
</thead>
<tbody>
<tr>
<td>DGCC(20:5/16:0)</td>
<td>+,-</td>
<td>1.00</td>
</tr>
<tr>
<td>LDGCC(20:5)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(22:6/16:0)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>LDGCC(22:6)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(20:5/20:5)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(22:6/20:5)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(20:5/16:1)</td>
<td>+,-</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(18:4/16:0)</td>
<td>+,-</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(16:1/16:0)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(20:5/14:0)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>LDGCC(18:4)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(16:1/16:1)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>TAG(16:0/16:0/14:0)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(20:5/18:2)</td>
<td>+</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(20:5/18:4)</td>
<td>+,-</td>
<td>1.00</td>
</tr>
<tr>
<td>Unknown</td>
<td>-</td>
<td>1.00</td>
</tr>
</tbody>
</table>

(B) Lipid ID             | Mode  | Normalised Abundance |
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>(Gly)$_2$Cer(d18:3/24:0)</td>
<td>-</td>
<td>1.00</td>
</tr>
<tr>
<td>LDGCC(16:1)</td>
<td>-</td>
<td>1.00</td>
</tr>
<tr>
<td>DGCC(18:2/16:0)</td>
<td>-</td>
<td>1.00</td>
</tr>
<tr>
<td>Unknown</td>
<td>-</td>
<td>0.86</td>
</tr>
<tr>
<td>SQDG(34:1)</td>
<td>-</td>
<td>0.78</td>
</tr>
<tr>
<td>Unknown</td>
<td>-</td>
<td>0.73</td>
</tr>
</tbody>
</table>
Fold Change (P-/P+)

<table>
<thead>
<tr>
<th>Lipid Identity</th>
<th>Fold Change</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cer(d18:3/24:0)</td>
<td></td>
</tr>
<tr>
<td>(Gly)_2Cer(d18:3/24:1)</td>
<td></td>
</tr>
<tr>
<td>(Gly)_2Cer(d18:3/24:0)</td>
<td></td>
</tr>
<tr>
<td>(Gly)_2Cer(d18:2/24:0)</td>
<td></td>
</tr>
<tr>
<td>(Gly)_2Cer(d18:1/24:0)</td>
<td></td>
</tr>
</tbody>
</table>