Genome-wide expression analysis of environmental stress in the cyanobacterium Synechocystis PCC 6803

Aguirre von Wobeser, E.

Publication date
2010

Link to publication

Citation for published version (APA):

General rights
It is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), other than for strictly personal, individual use, unless the work is under an open content license (like Creative Commons).

Disclaimer/Complaints regulations
If you believe that digital publication of certain material infringes any of your rights or (privacy) interests, please let the Library know, stating your reasons. In case of a legitimate complaint, the Library will make the material inaccessible and/or remove it from the website. Please Ask the Library: https://uba.uva.nl/en/contact, or a letter to: Library of the University of Amsterdam, Secretariat, Singel 425, 1012 WP Amsterdam, The Netherlands. You will be contacted as soon as possible.
Gene expression of the cyanobacterium *Synechocystis* PCC 6803 in response to nitrogen starvation

Vladimir Krasikov¹, Eneas Aguirre von Wobeser¹, Nataliya Yeremenko¹, Bas W. Ibelings², Jef Huisman¹ & Hans C. P. Matthijs¹,³

¹Aquatic Microbiology, Institute for Biodiversity and Ecosystem Dynamics, University of Amsterdam, Nieuwe Achtergracht 127, 1018 WS Amsterdam, the Netherlands

²Department of Foodweb Studies, Netherlands Institute of Ecology, Centre for Limnology, Nieuwersluis, the Netherlands

³Corresponding author: Hans Matthijs, phone: +31 20 5257070, fax: +31 20 5257064, Email: J.C.P.Matthijs@uva.nl
ABSTRACT

Acclimation to nitrogen starvation involves major changes in cyanobacterial physiology through regulation of gene expression. In this study, we report a genome-wide expression analysis of the response of the model cyanobacterium *Synechocystis* PCC 6803 after incubation of 12 h under nitrogen-free medium. For this purpose, we utilized an oligonucleotide microarray covering the whole genome of *Synechocystis* PCC 6803. Observed changes in gene expression were largely consistent with previous reports on short-term nitrogen starvation. The full transcriptome picture that was recorded after 12 h of nitrogen-free incubation showed that genes encoding components of the photosynthetic apparatus, including genes for phycobilisome synthesis, were significantly down-regulated. Conversely, phycobilisome degradation proteins were up-regulated. Furthermore, many genes that play key roles in nitrogen metabolism and transport of nitrogenous compounds were up-regulated. Nitrogen starvation caused a reduction of the transcription level of many genes related to carbon fixation. Genes encoding for carbon metabolism enzymes were a major point of regulation as well. We provide a novel interpretation of the carbon-related gene expression changes under nitrogen limitation, where carbon metabolism is prepared for rapid channelling of carbon reserves to nitrogen assimilation in case nitrogen would become available again. The results provide a detailed perspective of the functional changes in whole-cell metabolism of nitrogen-starved cyanobacteria.

INTRODUCTION

Low nitrogen availability limits the growth rate of cyanobacteria in a wide variety of freshwater and marine ecosystems (Goldman et al., 1990; Vitousek and Howarth, 1991). Most cyanobacteria use nitrate, nitrite or ammonium ions as nitrogen sources, with the preferred source being ammonium (Flores and Herrero, 2005). Some cyanobacteria may utilise urea, cyanate or amino acids, and some strains are also able to fix dinitrogen gas. Cyanobacteria have evolved sophisticated mechanisms to sense and respond to nitrogen limitation. Cyanobacteria typically bleach in response to prolonged nitrogen starvation; a phenomenon that is commonly known as chlorosis (Allen and Smith, 1969; Sauer et al., 2001). A number of genes and their products that participate in the process of chlorosis have been identified (Baier et al., 2001). A regulatory network driven by the transcription regulation factor NtcA (Herrero et al., 2001) and the regulatory protein PII (Hisbergues et al., 1999) has been studied in great detail. This regulatory network responds to the nitrogen status of the cell, perceived by sensing the intracellular concentration of 2-oxoglutarate, one of the major metabolites in the glutamine synthetase – glutamate synthase (GS-GOGAT) cycle responsible for ammonium assimilation (Muro-Pastor et al., 2001; Muro-Pastor et al., 2005).

During the past decade, sequence analyses of entire genomes have become available for several cyanobacteria, such as the freshwater cyanobacterium *Synechocystis* PCC 6803. Despite the relatively small genome size of *Synechocystis* (3.6 Mbp with 3264 ORFs), nearly half of its open reading frames (ORFs) still lack an assigned function. The availability of the genome sequence and annotation for *Synechocystis* PCC 6803 (Kaneko et al., 1996) enabled the construction of a DNA-microarray platform for global gene
expression analysis. The DNA-microarray approach has been successfully applied to study the whole-genome response of *Synechocystis* to high light and UV (Hihara *et al*., 2001; Huang *et al*., 2002; Tu *et al*., 2004), phosphate limitation (Suzuki *et al*., 2004), iron starvation-recovery (Singh *et al*., 2003), oxidative stress (Kobayashi *et al*., 2004), high salinity and osmotic shock (Kanesaki *et al*., 2002; Marin *et al*., 2003; Marin *et al*., 2004; Shoumskaya *et al*., 2005), cold-shock (Suzuki *et al*., 2001; Mikami *et al*., 2002), and to survey genes responsive in light-to-dark transitions (Gill *et al*., 2002).

The whole-genome response of cyanobacteria to nitrogen starvation was studied first for the nitrogen-fixing cyanobacterium *Anabaena* sp. PCC7120 (Ehira *et al*., 2003). This relatively early study made use of arrays that consisted of spotted fragments, each containing one to eight ORFs, rather than individually spotted genes. The study highlighted expressed sets of genes related to heterocyst formation and nitrogen assimilation. More recently, studies with oligonucleotide microarrays have focussed on different aspects of heterocyst formation in *Anabaena* sp. PCC 7120 (Ehira and Ohmori, 2006; Suzuki *et al*., 2006). Other recent studies have investigated the response of non-nitrogen-fixing cyanobacteria to nitrogen starvation (Krasikov *et al*., 2005; Osanai *et al*., 2006; Su *et al*., 2006). For instance, Osanai *et al.* (2006) presented the whole-genome response of *Synechocystis* PCC 6803 after an incubation of 4 h of nitrogen starvation. These authors found an induction of many sugar catabolic genes and a repression of sugar anabolic genes. They also showed the induction of several nitrogen related genes and the repression of photosynthetic genes and ribosomal protein genes. The persistence of these responses under long-term nitrogen starvation is so far unknown. Studies of the whole-genome responses of cyanobacteria to longer nitrogen starvation periods are therefore an exciting prospect in cyanobacterial genomics.

Here, we present the results of a genome-wide transcriptome analysis to further characterize the responses of *Synechocystis* PCC 6803 to nitrogen starvation. We have focussed on the response to nitrogen starvation occurring within one generation (12 h), a time span in which the acclimation process is already very active, but the observed physiological changes are not too dramatic yet.

**MATERIALS AND METHODS**

**Strains and growth conditions**

*Synechocystis* PCC 6803 was grown in 5 batch cultures at 30°C under continuous illumination (50 µmol photons m⁻² s⁻¹) in BG11 medium (Rippka *et al*., 1979). This nutrient-rich medium contains 17.6 mM inorganic nitrogen in the form of nitrate. All other nutrients are also at saturating concentrations in BG11 medium. Aeration was provided by CO₂-enriched air (3% on volume basis). Growth was monitored by measuring the optical density at 750 nm (OD₇₅₀) with a double-beam spectrophotometer (Aminco DW2000, USA). To induce nitrogen starvation, each batch culture of *Synechocystis* was grown to an optical density of OD₇₅₀ = 0.5 cm⁻¹, and subsequently divided into two sub-samples. Each sub-sample was harvested by centrifugation. Thereafter, one sub-sample of each culture was re-suspended in batch culture with the nitrogen-rich BG11 medium to an OD₇₅₀ of 0.1 cm⁻¹ and allowed to grow. These yielded 5 batch cultures, which were referred to as the
controls. The other set of sub-samples was re-suspended in batch culture with BG11\textsuperscript{0} medium lacking inorganic nitrogen, washed once in this medium by centrifugation, and next inoculated in fresh BG11\textsuperscript{0} to an OD\textsubscript{750} of 0.1 cm\textsuperscript{-1} and allowed to grow. This yielded 5 batch cultures, which were referred to as the nitrogen-starvation treatment. After 12 h of growth, all cultures were harvested by centrifugation. Cells pellets were frozen immediately in liquid nitrogen and stored at -80°C, for later RNA isolation.

RNA isolation and DNA microarray analysis

Total RNA was isolated by hot acid phenol and chloroform extraction, followed by LiCl precipitation and final RNA clean-up with a RNeasy mini kit (Qiagen, Germany). This method routinely removes soluble carbohydrates from RNA samples which improves the spectral characteristics of the RNA preparation. Quality and quantity of the obtained RNA were monitored using Nanodrop spectra (ND-1000, Nanodrop Technologies, USA) and Bioanalyzer assays (Agilent 2100 bioanalyzer, Agilent Technologies, USA). Single-stranded cDNA was prepared in the presence of Cy3-dUTP or Cy5-dUTP (Amersham Biosciences, UK) fluorescent dyes as follows. 10 µg total RNA and 0.5 µg of random hexamer (Amersham Biosciences) were mixed in total volume of 15 µL in water and the mixture was heated at 70°C for 10 min and then chilled on ice. The reverse transcription reaction was performed by addition to a final concentration of 10 mM DTT (Invitrogen, USA), 500 µM each of dATP, dCTP, and dGTP, 100 µM dTTP (dNTP set, Amersham Biosciences), and 65 µM Cy3-dUTP or Cy5-dUTP (Amersham Biosciences) in 30 µL of 1x First-Strand Buffer (Invitrogen). The mixture was incubated for another 10 min at 25°C. Subsequently, 400 units of Superscript II Reverse Transcriptase (Invitrogen) were added and the mixture was incubated for 110 min at 42°C. Next, the reaction was stopped and the RNA was hydrolyzed by addition of 1.5 µL of 1 M NaOH, followed by incubation for 10 min at 70°C. Thereafter, 1.5 µL of 1 M HCl was added for neutralization. Not-incorporated fluorescent nucleotides were removed with a QIAquick PCR purification kit (Qiagen). The reverse transcription and fluorescent dye incorporation efficiency was monitored with Nanodrop spectra. Cy3 and Cy5 labelled cDNA probes from control and nitrogen-starved cells, respectively, were combined, denatured for 2 min at 98°C, and mixed with Control Targets and 2x Hybridization Buffer (Agilent in situ hybridization kit-plus, Agilent Technologies). The mixture was then loaded on the glass slide. Hybridization and washing conditions were as described in the “Agilent 60-mer oligo microarray processing protocol, 6-screw chamber, SSPE Wash, version 2.0” (Agilent Technologies). Slides were scanned at 10 micron resolution in an Agilent microarray scanner and the spot intensities profiles thus derived were extracted by Feature Extraction Software version 7.5 (Agilent Technologies). To eliminate artefacts from differences in dye incorporation efficiency between the Cy3 and Cy5 dyes, the dye-swap strategy was applied (Churchill, 2002), and each sample was labelled with each of the dyes. To estimate the reliability of the microarray platform, self-self hybridizations have been performed. One microarray slide presented problems in the hybridization reaction, and was discarded, leaving 4 independent replicate cultures for the subsequent analysis.
Microarray design and data treatment

In order to investigate the transcriptional status of *Synechocystis*, we designed custom-made 45-65-mer oligonucleotides microarrays, which were printed for us by Agilent in 11K format on a standard 1x3 inch glass slide. Each of the 3264 genes of *Synechocystis* PCC 6803 (the full sequence and annotation is available at CyanoBase – http://www.kazusa.or.jp/cyano/) is represented in the array design by 1 to 4 different oligonucleotides dependent on the length of the ORF concerned. The complete design strategy for this oligonucleotide microarray for *Synechocystis* is described in Chapter 2 of this thesis.

Microarray data analysis

Statistical analysis of the microarray data was based on normalization of the data, followed by tests of the statistical significance of changes in gene expression, and finalized by a control step limiting the number of false positives.

Normalization of the microarray data was conducted without background subtraction, as has been recommended for Agilent microarrays (Zahurak *et al*., 2007). The data were log transformed, using natural logarithms, to reduce the skewness of the data distribution. To enable comparison of the data from all the microarray series, we utilized a parametric normalization procedure which was designed specifically for data sets with highly expressed genes (Chapter 3 of this thesis). This normalization procedure models the distribution of the microarray data using a Generalized Extreme Value (GEV) distribution, and subsequently converts the microarray data to a common distribution for all microarrays. The normalization procedure also adjusts partially or fully saturated values to make these values comparable between samples (for example, to compare saturated with down-regulated signals). After normalization, the 12 data-series had a common GEV distribution (see Chapter 3 for details).

Next, we proceeded to test for significant differences between the control treatments and the nitrogen-starvation treatments, using a gene by gene four-way Analysis of Variance (ANOVA). The gene by gene ANOVA model was modified from Kerr and Churchill (2001) and Wolfinger *et al*. (2001):

\[
 r_{gabjk} = G_g + (GT)_{ga} + P_b + (GTP)_{gab} + (GA)_{gj} + (GS)_{gk} + \epsilon_{gabjk}
\]

where \( r_{gabjk} \) is the normalized intensity for gene \( g \) (\( g=1,...,3264 \)), treatment \( a \) (\( a=0,1 \)), probe \( b \) (\( b=1,...,8091 \)), array \( j \) (\( j=1,...,6 \)) and sample \( k \) (\( k=1,...,6 \)), \( G_g \) is the main effect of each gene, \( GT_{ga} \) is the interaction effect between gene and treatment, \( P_b \) is the main effect of each probe, \( GTP_{gab} \) is the probe-specific treatment effect on each gene, \( GA_{gj} \) is the interaction effect between gene and array, and \( GS_{gk} \) is the interaction effect between gene and sample. The error term \( \epsilon_{gabjk} \) was assumed to be normally distributed with zero mean and variance \( \sigma^2 \). Note that the \( G \) term represents the fact that the model is fitted gene by gene. We included the \( GS \) term because the samples were randomized in the different arrays rather than hybridized in pairs, which allows quantification of the contributions of the different samples to the signal of each gene. The difference in gene expression between
the control treatment and the nitrogen-starvation treatment is determined by the values of the GT coefficients. Differences in gene expression between treatment and control samples hybridized to the same microarray are typically expressed as the log ratio (M) of the two signals. The GT coefficient is essentially equivalent to the log ratio, but takes into account all the microarrays of the experiment. For each of the 3264 genes of Synechocystis PCC 6803, the ANOVA produces a p-value indicating the significance of the GT term (i.e., the significance of changes in gene expression in response to nitrogen starvation).

The p-values obtained from the ANOVA estimate the probability of declaring a gene differentially expressed when it actually is not differentially expressed (error type 1). One needs to choose a critical p-value (p_c) below which all genes are considered significantly differentially expressed. Since we tested a total of 3264 hypothesis, we expected many false positives (i.e., many of the p-values might be low merely by chance). For instance, if one investigates 1000 genes, and declares genes as differentially expressed if p < 0.05, then one may expect 50 false positives. Therefore, we used the q-value method of Storey and Tibshirani (2001) to limit the false discovery rate (q), which is an estimate of the proportion of false positives among the genes declared significant. The false discovery rate is a function of p_c, and can be calculated as

$$ q(p_c) = \frac{N_f}{N_s} $$

where N_f is the expected number of false positives, and N_s is the number of genes declared significant (significant genes). We choose a critical p-value corresponding to a false discovery rate of q=0.01, to obtain an expected 1% of false positives among the significant genes.

RESULTS AND DISCUSSION

DNA microarray analysis

RNA from 4 nitrate-starved and 4 control Synechocystis PCC 6803 cultures was extracted, labelled and hybridized to 45-60-mer oligonucleotide microarray slides. The selected false discovery rate of q = 0.01 corresponded to a critical p-value of p < 0.0142. In total, 1485 genes (45% of the genome) were differentially expressed at a significance level of p < 0.0142 (Figure 1). Accordingly, the number of false positives was estimated to be about 15 genes. Of the 1485 genes with significantly different expression in the control and treatment samples, 835 genes were up-regulated under nitrogen starvation and 650 genes were down-regulated. Complete information of all significantly regulated genes is listed in Appendix 4A (up-regulated genes) and 4B (down-regulated genes). The results obtained were similar to the results reported by Osanai et al. (2006), since the M values of both studies were positively correlated with a correlation coefficient of ρ=0.7 (product-moment correlation: R²=0.5, n=3264, p<0.001), which is very high considering that the studies were performed in different labs and differed in the duration of nitrogen starvation (4 h vs. 12 h).
Figure 1. Response (M) of Synechocystis PCC 6803 gene expression to 12 h nitrogen starvation as a function of signal intensity (A) on the microarrays. The A values were calculated as the mean normalized signal intensity of all the probes for each gene on all arrays. The M values were estimated using analysis of variance as the gene by treatment interaction coefficients. Light gray points represent genes whose expression was not significantly altered by the treatment, whereas black dots represent the genes significantly responding to the treatment (p<0.0142, q=0.01). Values are in natural logarithmic scale.

Nitrogen uptake and assimilation genes

Increased efficiency of nitrogen assimilation is crucial for survival under nitrogen-depleted conditions. Accordingly, many genes involved in nitrogen uptake and nitrogen transport were found up-regulated after 12 h of nitrogen starvation (Table 1). In particular, the active uptake system for nitrate/nitrite (nrtABC) together with nitrite reductase (nirA), genes for glutamine permease (bgtB), urea transport genes (urtD and urtA) and cyanate lyase (cynS) were significantly up-regulated. The ammonium permease gene amt1 was significantly induced, but not amt2 and amt3. The gene amt1 is generally responsible for 95% of the ammonium uptake at very low ammonium concentrations (Montesinos et al., 1998), which are typical for natural habitats.

Intracellular ammonium, produced from the reduction of nitrate and nitrite, is incorporated into amino acids by sequential action of glutamine synthetase (GS) and glutamate synthase (GOGAT). Significant up-regulation was observed only for the GS type III that is encoded by the glnN gene (Table 1), suggesting that in particular the activity of this enzyme is required under nitrogen deficiency, as has been observed before (Reyes et al., 1997).
Table 1. Responses to nitrogen starvation of genes involved in nitrogen metabolism.

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change*</th>
<th>p-value</th>
<th>Gene function</th>
</tr>
</thead>
<tbody>
<tr>
<td>sll1423</td>
<td>ntcA</td>
<td>-0.7</td>
<td>0.04</td>
<td>global nitrogen regulator</td>
</tr>
<tr>
<td>sll0395</td>
<td>ntcB</td>
<td>1.1</td>
<td>0.82</td>
<td>nitrate assimilation transcriptional activator, LysR family protein</td>
</tr>
<tr>
<td>sso0707</td>
<td>glhB</td>
<td>2.4</td>
<td>9.03X10^{-11}</td>
<td>nitrogen regulatory protein P-II</td>
</tr>
<tr>
<td>sll1771</td>
<td>ppbA</td>
<td>1.1</td>
<td>0.48</td>
<td>protein serin-threonin phosphatase</td>
</tr>
<tr>
<td>slr2031</td>
<td>rsbU</td>
<td>1.1</td>
<td>0.89</td>
<td>putative PP2C-type protein phosphatase, required to recover from N or S starvation induced stationary phase</td>
</tr>
<tr>
<td>sso0452</td>
<td>nbla1</td>
<td>1.16</td>
<td>9.99X10^{-16}</td>
<td>phycobilisome degradation protein NblaA</td>
</tr>
<tr>
<td>sso0453</td>
<td>nbla2</td>
<td>25.8</td>
<td>2.61X10^{-4}</td>
<td>phycobilisome degradation protein NblaA</td>
</tr>
<tr>
<td>sll1151</td>
<td>glfB</td>
<td>-0.3</td>
<td>5.55X10^{-5}</td>
<td>glutamine synthetase inactivating factor IF17</td>
</tr>
<tr>
<td>sll1911</td>
<td>glfA</td>
<td>-0.4</td>
<td>0.02</td>
<td>glutamine synthetase inactivating factor IF7</td>
</tr>
<tr>
<td>sll0452</td>
<td>nblA1</td>
<td>16.6</td>
<td>9.99X10^{-16}</td>
<td>phycobilisome degradation protein NblA</td>
</tr>
<tr>
<td>sll0453</td>
<td>nblA2</td>
<td>25.8</td>
<td>2.61X10^{-4}</td>
<td>phycobilisome degradation protein NblA</td>
</tr>
</tbody>
</table>

**Nitrate/Nitrite uptake**

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change*</th>
<th>p-value</th>
<th>Gene function</th>
</tr>
</thead>
<tbody>
<tr>
<td>sll1450</td>
<td>nrtA</td>
<td>6.8</td>
<td>2.89X10^{-8}</td>
<td>nitrate/nitrite transport system substrate-binding protein</td>
</tr>
<tr>
<td>sll1451</td>
<td>nrtB</td>
<td>2.0</td>
<td>2.53X10^{-7}</td>
<td>nitrate/nitrite transport system permease protein</td>
</tr>
<tr>
<td>sll1452</td>
<td>nrtC</td>
<td>1.5</td>
<td>0.02</td>
<td>nitrate/nitrite transport system ATP-binding protein</td>
</tr>
<tr>
<td>sll1453</td>
<td>nrtD</td>
<td>1.1</td>
<td>0.58</td>
<td>nitrate/nitrite transport system ATP-binding protein</td>
</tr>
<tr>
<td>sll1454</td>
<td>nrtE</td>
<td>-0.9</td>
<td>0.34</td>
<td>ferredoxin-nitrate reductase</td>
</tr>
</tbody>
</table>

**Ammonium uptake**

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change*</th>
<th>p-value</th>
<th>Gene function</th>
</tr>
</thead>
<tbody>
<tr>
<td>sll0108</td>
<td>amt1</td>
<td>4.1</td>
<td>2.63X10^{-5}</td>
<td>ammonium/methylammonium permease</td>
</tr>
<tr>
<td>sll0537</td>
<td>amt3</td>
<td>1.2</td>
<td>0.31</td>
<td>ammonium/methylammonium permease</td>
</tr>
<tr>
<td>sll1017</td>
<td>amt2</td>
<td>1.7</td>
<td>0.18</td>
<td>ammonium/methylammonium permease</td>
</tr>
</tbody>
</table>

**Ammonium assimilation**

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change*</th>
<th>p-value</th>
<th>Gene function</th>
</tr>
</thead>
<tbody>
<tr>
<td>sll1289</td>
<td>icd</td>
<td>2.0</td>
<td>5.59X10^{-8}</td>
<td>isocitrate dehydrogenase (NADP+)</td>
</tr>
<tr>
<td>sll1756</td>
<td>gluN</td>
<td>1.2</td>
<td>0.42</td>
<td>glutamate–ammonia ligase</td>
</tr>
<tr>
<td>sll0288</td>
<td>glsN</td>
<td>2.2</td>
<td>1.93X10^{-4}</td>
<td>glutamate–ammonia ligase</td>
</tr>
<tr>
<td>sll1499</td>
<td>glsF</td>
<td>-0.8</td>
<td>0.22</td>
<td>ferredoxin-dependent glutamate synthase</td>
</tr>
<tr>
<td>sll1502</td>
<td>gltB</td>
<td>1.1</td>
<td>0.73</td>
<td>NADH-dependent glutamate synthase large subunit</td>
</tr>
<tr>
<td>sll1027</td>
<td>gltD</td>
<td>1.2</td>
<td>0.45</td>
<td>NADH-dependent glutamate synthase small subunit</td>
</tr>
<tr>
<td>sll1145</td>
<td>gltS</td>
<td>1.4</td>
<td>0.54</td>
<td>Monocomponent sodium-dependent glutamate permease GltS</td>
</tr>
<tr>
<td>sll0710</td>
<td>gdhA</td>
<td>-0.5</td>
<td>4.13X10^{-5}</td>
<td>glutamate dehydrogenase (NADP+)</td>
</tr>
<tr>
<td>sll1735</td>
<td>bgdA</td>
<td>1.2</td>
<td>0.62</td>
<td>ATP-binding subunit of the ABC-type Bgt permease for basic amino acids and glutamine</td>
</tr>
</tbody>
</table>

**Urea uptake**

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change*</th>
<th>p-value</th>
<th>Gene function</th>
</tr>
</thead>
<tbody>
<tr>
<td>sll0374</td>
<td>urtE</td>
<td>1.5</td>
<td>0.05</td>
<td>urea transport system ATP-binding protein</td>
</tr>
<tr>
<td>sll0764</td>
<td>urtD</td>
<td>2.7</td>
<td>2.99X10^{-11}</td>
<td>urea transport system ATP-binding protein</td>
</tr>
<tr>
<td>sll0447</td>
<td>urtA</td>
<td>2.9</td>
<td>1.78X10^{-4}</td>
<td>periplasmic protein, ABC-type urea transport system substrate-binding protein</td>
</tr>
<tr>
<td>sll1200</td>
<td>urtB</td>
<td>1.7</td>
<td>7.17X10^{-8}</td>
<td>urea transport system permease protein</td>
</tr>
<tr>
<td>sll1201</td>
<td>ureA</td>
<td>1.3</td>
<td>0.70</td>
<td>urea transport system permease protein</td>
</tr>
<tr>
<td>sll1256</td>
<td>ureB</td>
<td>1.1</td>
<td>0.23</td>
<td>urease gamma subunit</td>
</tr>
<tr>
<td>sll0420</td>
<td>ureC</td>
<td>1.3</td>
<td>0.12</td>
<td>urease beta subunit</td>
</tr>
<tr>
<td>sll1750</td>
<td>ureD</td>
<td>1.2</td>
<td>0.35</td>
<td>urease alpha subunit</td>
</tr>
<tr>
<td>sll1639</td>
<td>ureE</td>
<td>-0.7</td>
<td>2.65X10^{-4}</td>
<td>urease accessory protein D</td>
</tr>
<tr>
<td>sll1219</td>
<td>ureF</td>
<td>1.3</td>
<td>0.49</td>
<td>urease accessory protein E</td>
</tr>
<tr>
<td>sll1899</td>
<td>ureG</td>
<td>1.2</td>
<td>0.25</td>
<td>urease accessory protein G</td>
</tr>
</tbody>
</table>

**Cyanate uptake**

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change*</th>
<th>p-value</th>
<th>Gene function</th>
</tr>
</thead>
<tbody>
<tr>
<td>sll0899</td>
<td>cvnS</td>
<td>2.2</td>
<td>1.88X10^{-13}</td>
<td>cyanase</td>
</tr>
</tbody>
</table>

Notes: * Genes with positive fold changes were up-regulated in response to nitrogen starvation, while genes with negative fold changes were down-regulated. Genes with significant changes in gene expression (p<0.0142; q=0.01) are printed in bold.
Alternatively, ammonium can be incorporated directly into glutamate by glutamate dehydrogenase (gdhA), in a less efficient but less energy costly reaction. Concomitantly with the increased expression of glnN, the expression of gdhA was repressed (Table 1), showing the need for a high efficiency of this key reaction when energy is abundant. The supply of carbon skeletons for the GS-GOGAT cycle in the form of 2-oxoglutarate is provided by isocitrate dehydrogenase (encoded by the icd gene). The transcript level for this gene was significantly increased (Table 1), providing the GS-GOGAT cycle optimally with 2-oxoglutarate obtained from preceding carbon metabolism. Two small proteins, IF7 and IF17 (encoded by gifA and gifB genes), inhibiting the activity of GS by direct protein-protein interactions (García Domínguez et al., 1999), were strongly down-regulated (Table 1). These genes are in turn under the control of the global nitrogen regulator NtcA (García Domínguez et al., 2000). In our study, the ntcA gene did not show any significant change of expression level, though many regulated genes are under its direct transcriptional control, including cynS (Harano et al., 1997), icd (Muro-Pastor et al., 1996), and probably glnA, urtAB, nirA, glnB, amt1, and ndh (Su et al., 2005). Osanai et al. (2006) did not find expression changes of ntcA after 4 h of N starvation either. NtcA-mediated regulation of gene expression is controlled by complex interactions between this regulator and other molecules. For instance, ntcA interacts with the signal transduction protein PII through the mediation of pipX (Espinosa et al., 2006, 2007). The apparent lack of induction of ntcA expression suggests that the physiological state of NtcA might be more important for its regulation function than the concentration of NtcA. Alfonso et al. (2001) showed that the ntcA gene is transcribed at two different promoters, and one of the products is actually induced by nitrogen starvation. However, the change in the total ntcA mRNA concentration is apparently too low to be detected without the discrimination of the two transcripts, as is the case with microarrays. In contrast, the PII protein gene glnB, which can sense the nitrogen, carbon and energy balance of the cell by binding to 2-oxoglutarate and ATP (Forchhammer, 2004) was up-regulated under nitrogen starvation (Table 1). This is consistent with the study of Osanai et al. (2006), and previous reports (García Domínguez and Florencio, 1997).

Summarizing, we found gene expression changes in all the major components of nitrogen physiology. The changes consisted mostly of up-regulations, indicating an attempt by the cell to increase its nitrogen acquisition. However, down-regulation of three genes, gdhA and gifAB, may be as important to achieve that same purpose. Our results indicate that after 12 h of nitrogen starvation Synechocystis cells actively invest in the uptake of any nitrogen-containing compounds.

General stress response genes

The gene that presented the strongest up-regulation in this experiment was slr0376. Earlier research has shown that this gene with unknown function responds to various environmental stresses including nitrogen starvation (Singh and Sherman, 2002). The gene is co-transcribed as a 2.4 kb transcript with two other unknown genes, slr0373 and slr0374. These two genes ranked number 6 and 33, respectively, in our list of up-regulated genes in response to nitrogen starvation (Appendix 4A). The three genes are separated by long intergenic spacers. It is known that the full 2.4 kb transcript readily splits into smaller sub-transcripts, and that the relative abundances of these sub-transcripts vary among different stress conditions (Singh and Sherman, 2002). Differential expression of the three co-
transcribed genes therefore suggests differences in transcript stability after the initial transcription of DNA into polycistronic mRNA.

**Genes encoding terminal respiratory oxidases**

After 12 h of nitrogen starvation, there was an increased expression of all genes encoding terminal respiratory oxidases (Appendix 4A). The increased expression of terminal respiratory oxidases could be a means to dissipate reduction equivalents in the thylakoidal membranes, in order to protect photosystem II from photoinhibition.

Table 2. Number of photosynthesis and respiration genes significantly up-regulated or down-regulated after 12 h of nitrogen starvation (p<0.0142; q=0.01).

<table>
<thead>
<tr>
<th></th>
<th>Down-regulated</th>
<th>Up-regulated</th>
<th>Total number of genes*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Photosynthesis</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Photosystem I</td>
<td>16</td>
<td>0</td>
<td>16</td>
</tr>
<tr>
<td>Photosystem II</td>
<td>22</td>
<td>1</td>
<td>28</td>
</tr>
<tr>
<td>cytochrome b6/f</td>
<td>6</td>
<td>2</td>
<td>9</td>
</tr>
<tr>
<td>ATPase</td>
<td>10</td>
<td>0</td>
<td>10</td>
</tr>
<tr>
<td>NADPH dehydrogenase</td>
<td>6</td>
<td>4</td>
<td>22</td>
</tr>
<tr>
<td>Phycobilisomes</td>
<td>14</td>
<td>2</td>
<td>18</td>
</tr>
<tr>
<td>Soluble electron carriers</td>
<td>4</td>
<td>4</td>
<td>15</td>
</tr>
<tr>
<td><strong>Respiration</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Respiratory terminal oxidases</td>
<td>0</td>
<td>8</td>
<td>9</td>
</tr>
</tbody>
</table>

*Note: Total number of genes according to the gene categories defined by Cyanobase (http://www.kazusa.or.jp/cyano/).

**Genes involved in photosynthesis and carbon fixation**

We observed a major decrease in the expression of all components of the photosynthetic apparatus (Table 2). Almost all genes encoding subunits of Photosystems I and II, components of the phycobilisomes, and all subunits of ATP synthetase were down-regulated in response to nitrogen starvation. Several soluble electron transporters were down-regulated as well, probably anticipating a reduced demand for energy and reducing power in cellular metabolism. Nitrogen starvation leads to reduced growth rates (Richaud et al., 2001), which could be sustained with a reduced photosynthetic activity. Moreover, the photosynthetic apparatus contains a significant fraction of the total nitrogen of cyanobacterial cells under normal growth conditions (MacKenzie et al., 2005). Therefore, a reduction of the photosynthetic apparatus serves two purposes, to decrease the production of photosynthetic energy to match its reduced demand and to reduce allocations of nitrogen to the photosynthetic machinery. This partial arrest of photosynthesis precedes chlorosis (Sauer et al., 2001), which is typically observed later in time than the 12 h period of nitrogen starvation that we focussed on in this study. Chlorosis first involves the controlled degradation of the phycobilisomes and thereafter of the photosynthetic machinery as a whole. Our results show that genes for two small phycobilisome-degradation proteins,
Figure 2. Up- and down-regulation of carbon metabolism gene expression of *Synechocystis* PCC 6803 after 12 h nitrogen starvation displayed on a metabolic pathway map. The pathway map was adapted from Osanai *et al.* (2007) and the Kyoto Encyclopedia of Genes and Genomes (http://www.genome.jp/kegg/). Red arrows indicate up-regulation and blue arrows down-regulation. For the gene descriptions and differential expression data, see Table 3. ADP-Glc is ADP-glucose; G1P, glucose-1-phosphate; G6P, glucose-6-phosphate; 6PGL, 6-phosphogluconolactone; 6PG, 6-phosphogluconate; RU5P, ribulose-5-phosphate; X5P, xylulose-5-phosphate; R5P, ribose-5-phosphate; E4P, erythrose-4-phosphate; S7P, sedoheptulose-7-phosphate; RU1.5P, ribulose-1,5-biphosphate; F6P, fructose-6-phosphate; F1,6P2, fructose-1,6-biphosphate; DHAP, dihydroxyacetone phosphate; GAP, glyceraldehyde-3-phosphate; G1,3P2, 1,3-bisphosphoglycerate; G3P, 3-phosphoglycerate; G2P, 2-phosphoglycerate; PEP, phosphoenolpyruvate; PYR, pyruvate; ACA, acetyl coenzyme A; OAA, oxaloacetate; CIT, citrate; ICI, isocitrate; OXO, 2-oxoglutarate.

*NblA1* and *NblA2*, increased their transcript level more than tenfold (Table 1). Li and Sherman (2002) showed a high accumulation of nblA mRNA after a 24 h long nitrogen starvation period. In fact, together with the previously gene encoding a protein of unknown function (*slr0376*), these two phycobilisome-degradation proteins are placed in the top three of the most strongly up-regulated genes in response to nitrogen starvation (Appendix 4A). The presence of these genes is known to be essential for directed phycobilisome
degradation (Baier et al., 2001; Richaud et al., 2001), although the actual proteases involved in this process have not yet been characterized.

Consistent with the reduced expression of many genes of the photosynthetic apparatus, the pathway for photosynthetic carbon fixation, the Calvin cycle, also had many of its genes down-regulated under nitrogen starvation (Table 3; Figure 2). Particularly, the genes encoding Ribulose-1,5-bisphosphate carboxylase/oxygenase (RUBISCO), which represents an important nitrogen pool in nitrogen replete cells (MacKenzie et al., 2005), were strongly down-regulated, as has been observed before (Aldehni et al. 2003; Osanai et al., 2006). These observations indicate a sharp decline in the demand for carbon skeletons in response to nitrogen starvation. This is consistent with the further observation that genes encoding several components of the carboxysomes and their carbon concentration mechanism proteins were also down-regulated, which could lead to a decreased efficiency of RUBISCO, whose activity depends on a CO₂-rich microenvironment created by these proteins.

Genes involved in central carbon metabolism

The central carbon metabolism of Synechocystis PCC 6803 was extensively regulated at the gene expression level after 12 h of nitrogen starvation (Table 3). Figure 2 presents the up-regulated and down-regulated genes of the central carbon metabolism of Synechocystis PCC 6803 after 12 h of nitrogen starvation. This metabolic map assumes that all metabolites involved are accessible to all enzymes at all times (i.e., the no compartmentation assumption). Our results confirm the finding of Osanai et al. (2006) that gene expression of sugar catabolic enzymes is induced under nitrogen starvation, while genes involved in sugar anabolism are repressed (Table 3, Figure 2). While genes acting exclusively in the direction of gluconeogenesis were down-regulated (fbp and gapII), genes acting exclusively on the glycolytic path were induced (pfk and gap1). This is in apparent contradiction with the clearly established accumulation of glycogen, a product of sugar anabolism, under nitrogen starvation (Tandeau de Marsac et al., 1980; Osanai et al., 2006). Osanai et al. (2006) point out that glgB, a gene involved in the synthesis of glycogen, was induced after 4 h nitrogen starvation, an observation confirmed by our results after 12 h nitrogen starvation. However, the induction of glgB is not enough to explain the accumulation of glycogen, since several enzymes acting before glgB on the glycogen synthesis pathway were down-regulated. Rather, the flow of carbon from the Calvin cycle to glycogen synthesis needs to match the degradation of glycogen in order to maintain a large glycogen pool. Metabolic fluxes cannot be predicted with gene expression data alone (Yang et al., 2002), since they depend strongly on the concentrations of the substrates and products of the individual reactions, as well as on allosteric regulation of enzyme activities. We propose that carbon fixation, although itself repressed at the gene expression level (Table 2, Figure 3), is high enough to push the chemical equilibrium towards sugar anabolism, leading to glycogen accumulation.

The accumulation of 2-oxoglutarate under nitrogen starvation, another well established fact (Muro-Pastor et al., 2001; Espinosa et al., 2006), implies carbon flux in the opposite, catabolic direction. We propose a branching point at 3-phosphoglycerate, where part of this metabolite would follow the sugar anabolic route, following the Calvin cycle/gluconeogenesis, and part would follow the glycolytic route and the tricarboxilic acid
cycle to produce 2-oxoglutarate for ammonium assimilation. The stoichiometry of the Calvin cycle implies that for every 3-phosphoglycerate molecule channeled to ammonium assimilation, 5 molecules would need to return to the Calvin cycle to regenerate the ribulose-5-phosphate pool. Additionally, several 3-phosphoglycerate molecules would need to follow this route to compensate for the use of other Calvin cycle compounds for other biosynthetic needs, like the use of 5-carbon sugars for nucleotide synthesis. Once the concentrations of the metabolites in the system are in equilibrium, the carbon flow to 2-oxoglutarate is probably controlled by the ammonium assimilation rate, limited by the nitrogen availability. The high concentration of 2-oxoglutarate probably makes its synthesis thermodynamically less favourable, and only possible at a rate similar to its consumption.

If more nitrogen would become available, more 3-phosphoglutamate would be pulled towards the Citric Acid cycle and ammonium assimilation, because the consumption of 2-oxoglutarate would favour its synthesis. The carbon skeletons driven away from the central intermediary metabolism would then probably be restituted from the glycogen pool. This would explain the up-regulation of the oxidative part of the pentose-phosphate cycle, which would allow the conversion of glucose-6-phosphate directly into ribulose-5-phosphate, with the generation of NADPH. With a sudden increase of nitrogen supply, the glycolytic pathway could also channel carbon from glycogen to 3-phosphoglutamate until the flow from carbon fixation pushes the reaction in the gluconeogenesis direction again. Under this scheme, the glycogen reserves would act as a buffer to compensate for changes in the carbon demand for ammonium assimilation (and other biosynthetic needs). This could ensure constant levels of the Calvin cycle intermediates, anticipating a rapid increase of Calvin cycle activity when nitrogen would become available again.

### Table 3. Responses to nitrogen starvation of genes involved in carbon metabolism. *

<table>
<thead>
<tr>
<th>ORF</th>
<th>Gene</th>
<th>Fold change</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Calvin cycle</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll1028</td>
<td>ccmK2</td>
<td>-3.58772</td>
<td>&lt;1X10^-16</td>
</tr>
<tr>
<td>sll1029</td>
<td>ccmK1</td>
<td>-3.23913</td>
<td>&lt;1X10^-16</td>
</tr>
<tr>
<td>sll1039</td>
<td>ccmK4</td>
<td>-2.55852</td>
<td>&lt;1X10^-16</td>
</tr>
<tr>
<td>sll0009</td>
<td>rbcL</td>
<td>-5.84948</td>
<td>7.77X10^-16</td>
</tr>
<tr>
<td>sll1031</td>
<td>ccmM</td>
<td>-2.4129</td>
<td>1.65X10^-13</td>
</tr>
<tr>
<td>sll0012</td>
<td>rbcS</td>
<td>-5.36463</td>
<td>1.5X10^-12</td>
</tr>
<tr>
<td>sll1388</td>
<td>ccmK3</td>
<td>-1.85192</td>
<td>1.67X10^-11</td>
</tr>
<tr>
<td>sll1347</td>
<td>icfA</td>
<td>-1.41546</td>
<td>6.45X10^-9</td>
</tr>
<tr>
<td>sll1525</td>
<td>prk</td>
<td>-2.41091</td>
<td>0.000119</td>
</tr>
<tr>
<td>sll0051</td>
<td>ecaB</td>
<td>1.262831</td>
<td>0.000185</td>
</tr>
<tr>
<td>sll1030</td>
<td>ccmL</td>
<td>-1.41535</td>
<td>0.037607</td>
</tr>
<tr>
<td>sll0436</td>
<td>ccmO</td>
<td>1.181517</td>
<td>0.155448</td>
</tr>
<tr>
<td>sll0934</td>
<td>ccmA</td>
<td>1.069244</td>
<td>0.237757</td>
</tr>
<tr>
<td><strong>Calvin cycle and glycolysis/gluconeogenesis</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll0394</td>
<td>pgk</td>
<td>-2.16666</td>
<td>3.67X10^-13</td>
</tr>
<tr>
<td>sll0204</td>
<td>fbpI</td>
<td>-2.20979</td>
<td>4.22X10^-11</td>
</tr>
<tr>
<td>sll0018</td>
<td>cbbA</td>
<td>-1.35063</td>
<td>1.08X10^-8</td>
</tr>
<tr>
<td>sll1342</td>
<td>gap2</td>
<td>-1.58593</td>
<td>1.84X10^-4</td>
</tr>
<tr>
<td>sll0783</td>
<td>tpi</td>
<td>1.260804</td>
<td>0.0001288</td>
</tr>
<tr>
<td>sll0952</td>
<td>fbpII</td>
<td>1.175966</td>
<td>0.048053</td>
</tr>
<tr>
<td>sll0943</td>
<td>cbbA</td>
<td>-1.04935</td>
<td>0.693025</td>
</tr>
</tbody>
</table>

---

*Calvin cycle: sll1028, ccmK2 -3.58772, <1X10^-16; sll1029, ccmK1 -3.23913, <1X10^-16; sll1039, ccmK4 -2.55852, <1X10^-16; sll0009, rbcL -5.84948, 7.77X10^-16; sll1031, ccmM -2.4129, 1.65X10^-13; sll0012, rbcS -5.36463, 1.5X10^-12; sll1388, ccmK3 -1.85192, 1.67X10^-11; sll1347, icfA -1.41546, 6.45X10^-9; sll1525, prk -2.41091, 0.000119; sll0051, ecaB 1.262831, 0.000185; sll1030, ccmL -1.41535, 0.037607; sll0436, ccmO 1.181517, 0.155448; sll0934, ccmA 1.069244, 0.237757.

**Calvin cycle and glycolysis/gluconeogenesis:** sll0394, pgk -2.16666, 3.67X10^-13; sll0204, fbpI -2.20979, 4.22X10^-11; sll0018, cbbA -1.35063, 1.08X10^-8; sll1342, gap2 -1.58593, 1.84X10^-4; sll0783, tpi 1.260804, 0.0001288; sll0952, fbpII 1.175966, 0.048053; sll0943, cbbA -1.04935, 0.693025. 
<table>
<thead>
<tr>
<th>Gene Symbol</th>
<th>Description</th>
<th>Expression Value</th>
<th>p-Value</th>
<th>q-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>slr1945</td>
<td>pgm -1.44804 9.99X10(^{-11}) 2,3-bisphosphoglycerate-independent phosphoglycerate mutase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0301</td>
<td>ppsA -1.84634 8.22X10(^{-11}) phosphoenolpyruvate synthase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0886</td>
<td>gap1 1.92705 1.52X10(^{-10}) glyceraldehyde 3-phosphate dehydrogenase 1 (NAD+)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1841</td>
<td>pyk 1.68134 2.84X10(^{-8}) pyruvate kinase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0301</td>
<td>ppsA -1.84634 8.22X10(^{-11}) phosphoenolpyruvate synthase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0886</td>
<td>gap1 1.92705 1.52X10(^{-10}) glyceraldehyde 3-phosphate dehydrogenase 1 (NAD+)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1841</td>
<td>pyk 1.68134 2.84X10(^{-8}) pyruvate kinase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll0587</td>
<td>odhB - 4.05X10(^{-8}) pyruvate dehydrogenase dihydrolipoamide acetyltransferase component (E2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1721</td>
<td>pdB - 3.27X10(^{-7}) pyruvate dehydrogenase E1 component, beta subunit</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0290</td>
<td>ppc 1.43632 4.16X10(^{-7}) phosphoenolpyruvate carboxylase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1196</td>
<td>pfkA 1.271872 3.39X10(^{-6}) phosphofructokinase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0752</td>
<td>eno 1.123512 0.00011 enolase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1124</td>
<td>gpmB -1.48576 0.000251 phosphoglycerate mutase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1275</td>
<td>pyk2 -1.19337 0.000551 pyruvate kinase 2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1349</td>
<td>pgI 1.126223 0.002693 glucose-6-phosphate isomerase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll0593</td>
<td>glk -1.27304 0.006133 glucone kinase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1096</td>
<td>phdD -1.13330 0.002031 pyruvate dehydrogenase dihydrolipoamide dehydrogenase (E3)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0745</td>
<td>pfkA 1.30439 0.054343 phosphofructokinase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll1275</td>
<td>pyk2 -1.19337 0.000551 pyruvate kinase 2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1349</td>
<td>pgI 1.126223 0.002693 glucose-6-phosphate isomerase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1934</td>
<td>-1.0852 0.59 pyruvate dehydrogenase E1 component, alpha subunit</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0214</td>
<td>zwf 1.394768 2.77X10(^{-9}) glucose 6-phosphate dehydrogenase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll0329</td>
<td>gnd 1.631945 4.38X10(^{-9}) 6-phosphogluconate dehydrogenase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1479</td>
<td>pgl 1.179716 0.000575 6-phosphogluconolactonase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1734</td>
<td>opeA 1.519312 0.001734 glucose 6-phosphate dehydrogenase assembly protein</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1793</td>
<td>tal 1.269939 0.252354 transaldolase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0665</td>
<td>acnB 1.805104 2.22X10(^{-16}) aconitate hydratase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll0870</td>
<td>rpe -1.63834 1.37X10(^{-9}) pentose-5-phosphate-5-epimerase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ss2153</td>
<td>rpiB 1.041954 0.374137 probable ribose phosphate isomerase B</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0194</td>
<td>rpiA 1.084116 0.538663 ribose 5-phosphate isomerase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1843</td>
<td>zwf 1.394768 2.77X10(^{-9}) glucose 6-phosphate dehydrogenase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sll0329</td>
<td>gnd 1.631945 4.38X10(^{-9}) 6-phosphogluconate dehydrogenase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1479</td>
<td>pgl 1.179716 0.000575 6-phosphogluconolactonase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1734</td>
<td>opeA 1.519312 0.001734 glucose 6-phosphate dehydrogenase assembly protein</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1793</td>
<td>tal 1.269939 0.252354 transaldolase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0665</td>
<td>acnB 1.805104 2.22X10(^{-16}) aconitate hydratase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1229</td>
<td>icd 1.950823 2.91X10(^{-9}) isocitrate dehydrogenase (NADP+)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0401</td>
<td>gtaA 1.360651 3.08X10(^{-9}) citrate synthase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1233</td>
<td>frdA 1.310954 1.5X10(^{-9}) succinate dehydrogenase flavoprotein subunit</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0891</td>
<td>citH, ldh -1.25475 0.00033 malate dehydrogenase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1157</td>
<td>sucD -1.18632 0.000864 succinyl-CoA synthetase alpha chain</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0018</td>
<td>fumC 1.188981 0.024007 fumarase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1123</td>
<td>sacB 1.154418 0.025585 probable succinate dehydrogenase iron-sulfur protein</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1162</td>
<td>sacB 1.044988 0.076864 succinate dehydrogenase iron- sulphur protein subunit</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1123</td>
<td>sacC -1.04068 0.313163 succinyl-CoA synthetase beta chain</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1393</td>
<td>glgA -1.24565 4.52X10(^{-5}) glycogen (starch) synthase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0945</td>
<td>glgA -1.541 0.000567 glycogen synthase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr0158</td>
<td>glgB 1.519001 1.05X10(^{-7}) 1.4-alpha-glucan branching enzyme</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1176</td>
<td>glgC -2.68343 1.82X10(^{-14}) glucose-1-phosphate dehydrogenase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1348</td>
<td>glgP 1.087603 7.42X10(^{-11}) glycogen phosphorylase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1356</td>
<td>glgP 1.607764 0.00199 glycogen phosphorylase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1237</td>
<td>glgX 1.580152 4.12X10(^{-5}) glycogen operon protein GlgX homolog</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1857</td>
<td>glgX 1.187306 1.68X10(^{-5}) isomylase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1676</td>
<td>malQ 1.359493 1.87X10(^{-5}) 4-alpha-glucanotransfer</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>slr1726</td>
<td>pgm -1.14825 0.013064 phsophoglucomutase</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Note: Genes with significant changes in gene expression (p<0.0142; q=0.01) are printed in bold.
In summary, the accumulation of 2-oxoglutarate and glycogen under nitrogen starvation suggests that the gene expression changes in central carbon metabolism are preparations for a fast response to increases in nitrogen availability. As an exception, down-regulation of the Calvin cycle is probably exerted to match the reduced photosynthetic activity and reduced demand for carbon skeletons for ammonium assimilation during nitrogen starvation.

CONCLUSION

In conclusion, nitrogen starvation triggers a general increase of expression of genes involved in nitrogen uptake and assimilation, reflecting an effort of the cells to increase the acquisition of nitrogen. Our results also show an increase in the enzymes leading to the production of 2-oxoglutarate, which probably assures the supply of carbon skeletons for ammonium assimilation, and maintains the nitrogen sensory systems based on the concentration of 2-oxoglutarate in an active state. Our results clearly demonstrate down-regulation of genes encoding proteins that are involved in photosynthesis, cell growth, and the synthesis of nitrogen-rich biomolecules. The overall protein expression levels were apparently reduced, since genes encoding all 53 ribosomal proteins and several constitutive subunits of RNA polymerase were down-regulated (Table S2). This is an obvious prelude to the resting state that cells may reach after prolonged nitrogen starvation (Sauer et al., 2001). The expression changes observed in genes involved in carbon metabolism suggest an anticipation of an increase in the nitrogen availability by an increased expression of genes for the degradation of glycogen, to support a constant carbon supply for ammonium assimilation and carbon fixation. We also found many examples of genes encoding unknown and hypothetical proteins that were either up- or down-regulated in response to nitrogen starvation (Appendix 4A and 4B), reflecting the fact that about 50% of the genes in the Synechocystis genome still lack a functional annotation. These unknown and hypothetical proteins offer a source of curiosity, where the construction of deletion mutants may help to render new insight into the molecular biology and physiology of cyanobacteria.

ACKNOWLEDGEMENTS

The research of VK, EAvW, and JH was supported by the Earth and Life Sciences Foundation (ALW), which is subsidized by the Netherlands Organization for Scientific Research (NWO). EAvW was further supported by the Consejo Nacional de Ciencia y Tecnología (CONACYT) of Mexico. BWI was supported by a grant of the Royal Dutch Academy of Sciences (KNAW). We kindly thank Dr. Floyd Wittink of the Microarray Department of the Faculty of Science of the University of Amsterdam for helpful advices and access to their bioanalyser and reader facilities.
REFERENCES


