A Tribute to Disorder in the Genome of the Bloom-Forming Freshwater Cyanobacterium Microcystis aeruginosa


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A Tribute to Disorder in the Genome of the Bloom-Forming Freshwater Cyanobacterium *Microcystis aeruginosa*

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Abstract

*Microcystis aeruginosa* is one of the most common bloom-forming cyanobacteria in freshwater ecosystems worldwide. This species produces numerous secondary metabolites, including microcystins, which are harmful to human health. We sequenced the genomes of ten strains of *M. aeruginosa* in order to explore the genomic basis of their ability to occupy varied environments and proliferate. Our findings show that *M. aeruginosa* genomes are characterized by having a large open pangenome, and that each genome contains similar proportions of core and flexible genes. By comparing the GC content of each gene to the mean value of the whole genome, we estimated that in each genome, around 11% of the genes seem to result from recent horizontal gene transfer events. Moreover, several large gene clusters resulting from HGT (up to 19 kb) have been found, illustrating the ability of this species to integrate such large DNA molecules. It appeared also that all *M. aeruginosa* displays a large genomic plasticity, which is characterized by a high proportion of repeat sequences and by low synteny values between the strains. Finally, we identified 13 secondary metabolite gene clusters, including three new putative clusters. When comparing the genomes of *Microcystis* and *Prochlorococcus*, one of the dominant picocyanobacteria living in marine ecosystems, our findings show that they are characterized by having almost opposite evolutionary strategies, both of which have led to ecological success in their respective environments.

Introduction

Cyanobacteria play an important role in aquatic ecosystems (e.g. [1,2]) but they can also disrupt the functioning of these ecosystems and their use by humans, because of the ability of several species to proliferate and to produce harmful toxins (e.g. [3]). In recent years, several studies have shown that sharply contrasting ecological strategies have allowed cyanobacteria to live in a large range of habitats in the euphotic zone of aquatic ecosystems. For example, in *Prochlorococcus marinus*, one of the dominant picocyanobacterial genera in marine ecosystems, different ecotypes occupy various ecological niches mainly defined by the availability of nutrients and light [4–7]. In contrast, in *Microcystis aeruginosa*, one of the most common toxic bloom-forming species in eutrophic freshwater ecosystems, no evidence of ecotype differentiation has been found. Moreover, population genetic studies have revealed that *M. aeruginosa* populations are characterized by wide genetic diversity and the lack of biogeographical patterns of genetic differentiation, suggesting that *M. aeruginosa* strains are able to proliferate in a wide range of ecosystems [8,9].

Hence, one exciting scientific challenge is to elucidate the genomic basis of the different ecological strategies developed by the *Prochlorococcus* and *Microcystis* genera. With this goal, the genomes of many *P. marinus* strains have recently been sequenced (for a review, see [10]). These studies have indicated that almost all sequenced *P. marinus* genomes are characterized by a small size that seems to result from a genome reduction process. However, despite their small size, these genomes have acquired a large number of genes by horizontal gene transfer (HGT) [11–15]. Thus, the combination of a niche specialization process involving genome reduction [16,17], and a high adaptive potential with gene acquisition by HGT, might explain the ecological success of this species.

In contrast to *Prochlorococcus*, only two *M. aeruginosa* strains have been sequenced so far. From these first two studies, *M. aeruginosa* genomes appear to display unusual plasticity, reflected in a large number of repeated sequences and low synteny values between...
them [18,19], but these findings should be confirmed on a greater number of strains. In addition, data are needed about the size of the core and pangenome and on the impact of HGT events on the ability of members of this species to colonize various environments. In this goal, we sequenced the genomes of ten other *M. aeruginosa* strains, selected on their phylogenetic relatedness, their geographical origins, and their ability to produce various secondary metabolites.

**Materials and Methods**

*M. aeruginosa* strain culture and DNA isolation

Axenic strains of *M. aeruginosa* were from the Pasteur Culture collection of Cyanobacteria (PCC) and from the NIES collection (T1-4). Cells were grown at 25°C in BG110 supplemented with 2 mM NaNO₃ and 10 mM NaHCO₃ [20]. They were harvested by centrifugation (10,000 g, 10 min, 18°C), washed twice with sterile distilled water, and kept frozen until DNA extraction.

DNA extraction from the frozen pellets was carried out using Genomic DNA isolation-NucleoBond ® AX (Macherey-Nagel, Hoerdt, France) according to the manufacturer’s instructions.

Sequencing and assembling methods

The sequences of the ten *M. aeruginosa* genomes were obtained by combining several approaches. First, a single read library was constructed for each of the ten *M. aeruginosa* strains and sequenced with the GSflex version (250 nt length), until 16 to 25-fold coverage was obtained. Then, 3 to 8-fold coverage 454 Titanium reads (average 450 nt length), obtained from mate-paired libraries with 2–5 kb or 6–8 kb insert sizes, were added. In agreement with other de novo sequencing projects, the percentage of 454 reads used for assembly was approximately 97% and the percentage of Illumina reads used for error correction around 93%. The assembly was performed using Newbler Software (v2.3, Roche). All sequences, which were not included in scaffolds after the assembly process and which displayed a

Illumina reads used for error correction around 93%. The for assembly was approximately 97% and the percentage of Illumina reads used for error correction around 93%. The assembly was performed using Newbler Software (v2.3, Roche).

Both coding sequence prediction and automatic annotation were performed by using the Microscope platform, a web-based framework for the systematic and efficient revision of microbial genome annotation (http://www.genoscope.cns.fr/arc/ microcope) [22]. Expert validations were carried out for specific genes. This platform was also used for the comparative genome analyses performed on the twelve *Microcystis* genomes and on those of other cyanobacterial species.

The overall characteristics of the ten *M. aeruginosa* genomes are reported in Table S1. Despite high genome coverage values, from 27 to 121 scaffolds were obtained per strain at the end of the assembly. The number of scaffolds was positively correlated to the number of other cyanobacterial species.

**Phylogenetic analyses**

We used two datasets to reconstruct phylogenetic relationships between twelve *M. aeruginosa* strains (the ten newly sequenced ones and two publicly available ones, PCC 7806 and NIES-843): (i) seven housekeeping genes used by Tanabe et al. [23], and (ii) 1989 genes from the core genome of the twelve *M. aeruginosa* strains. These 1989 genes among the 2462 core genes correspond to the orthologous genes displaying at least 80% sequence identity over at least 80% of the length of the smallest protein. Before concatenation, the homologous sequences of each gene were aligned using the MUSCLE software with default parameters [24] and the alignments were filtered by using the program GBLOCKS allowing half gap positions [23].

For each dataset, we computed trees using PHYML [26] and we used the Jones-Taylor-Thornton model of amino acid substitution for the protein dataset (core genes), and the Gamma Time Reversible model for the nucleic dataset (housekeeping genes). Heterogeneities between sites were estimated using gamma-distributed rate variation (4 categories), the alpha parameter was computed using PHYML, tree topologies were explored using Nearest Neighbor Interchanges and we used the tlr option to optimise the topology, the branch lengths and rate parameters of the starting trees. One hundred bootstrap replicates were performed to assess the statistical support of each node.

**Bioinformatic analyses**

All the bioinformatics analyses were performed by using tools provided by the MicroScope platform [22].

The orthoMCL program (version 1.4), which uses a Markov Cluster algorithm, was used to compute the core and the pangenome of *M. aeruginosa*. Putative orthologies were defined as gene pairs satisfying an alignment threshold of at least 50% amino acid sequence identity over at least 50% of the length of the smallest protein.

The proportion of repeats was estimated using the Repseek algorithm. This algorithm is a fast two-step method (seed detection followed by their extensions), which allows finding large degenerate repeats within or between large DNA sequences [27].

The synteny values representing the percentage of CDSs belonging to a syntenic group were estimated by taking into account CDSs sharing at least 35% sequence identity on 80% of the length of the smallest protein, with a gap parameter (number of consecutive genes not involved in synteny), which was set to five.

Finally, the evaluation of horizontal gene transfers was performed (i) by estimating the proportion of genes displaying more than 20% difference in their GC content compared to the mean GC content of the whole genome, and (ii) by the implementation of Interpolated Variable Order Motifs (IVOMs), which exploits compositional biases using variable order motif distributions (2mer to 8mer). This implementation was achieved by using the Alien-Hunter application included in the MicroScope platform.

**Secondary metabolites gene cluster identification**

A modified version of the complete genome scanning pipeline for searching secondary metabolites (http://nrps.igs.umaryland.edu/nrps/2metdb/) implemented on the Microscope platform was used to detect NRPS/PKS genes [28]. Each gene within a cluster was compared to its syntenic counterpart at the amino-acid level in the reference genome to obtain the deduced amino-acid sequence identity. The genome of the *M. aeruginosa* strain PCC 7806 [19] was used as reference but when the compared gene cluster was absent from this genome, those of PCC 7941, PCC 9432, PCC 9806 and NIES-843 [18] or the gene cluster psm3 from *Microcystis* sp. K139 [29] were used. Adenylation specificity was checked using online NRPS predictor (http://www-ab.informatik.uni-tuebingen.de/software/NRSPredictor; [30]).

**Results**

**General features of the ten *M. aeruginosa* genomes**

The size of the ten new *M. aeruginosa* genomes ranged from 4.2 to 5.2 Mbp. The GC-content value was around 43% for each of the genomes, and the Coding DNA Sequence (CDS) density of
found in two copies in the two strains belonging to the sub-clade 1 strains, as for example one of the ten new genomes (Table S3). A restricted number of 205 genes composing this Minimal Gene Set, none of the genes Phylogenetic studies of the twelve M. aeruginosa genomes described in Table 1 (data not shown), the NIES-843 genome was clearly distinguishable from all the other genomes. Finally, a comparison of the composition of the Minimal Gene Set [31] was performed on all the M. aeruginosa genomes with the goal to evaluate the quality of the ten new sequenced ones. Among the 205 genes composing this Minimal Gene Set, none of the genes found in NIES-843 [10] and in PCC7806 [19] was lacking in only one of the ten new genomes (Table S2). A restricted number of them displayed a heterogeneous distribution among the twelve strains, as for example pgi (phosphoglucone isomerase), which was found in two copies in the two strains belonging to the sub-clade 1 (see bellow), and only in one copy in the other ones.

Estimation of the core and pangenome in M. aeruginosa

As shown in Fig. 1B, the size of the core genome is close to 2462 genes (Table S4). As this curve is not completely asymptotic, the addition of new genomes would probably reduce slightly the size of the M. aeruginosa core genome. With regard to the pangenome, the gene accumulation curve does not reach a plateau, indicating that its size (>12000 genes) is largely underestimated (Fig. 1A).

When looking at distribution of the core and flexible genes classified among the Clusters of Orthologous Groups (COGs) categories, it appeared that a smaller proportion of flexible genes was found in C (Energy production and conversion), F (Nucleotide transport and metabolism), H (Coenzyme metabolism), I (Translation, ribosomal structure and biogenesis) and M (Cell envelope biogenesis, outer membrane) categories, compared to that of core genes. In contrast, this proportion of flexible genes is high in the L category (DNA replication, recombination and repair) due to the presence of numerous transposase encoding genes, and also in R (General function prediction only) and S (Unknown function) categories (Fig. 2; Table S4).

Phylogenetic studies of the twelve M. aeruginosa strains and relationships with their geographical origin and their genome size

A phylogenetic analysis was performed on seven housekeeping genes (ftsZ, glaA, glnA, gyrB, pgi, recA and tpi) and on 1989 genes belonging to the core genome. The two trees were congruent, but phylogenetic analysis based on the core genome enhanced the differentiation of subclades (SC) 3 and 4 (Fig. 3). These two trees were also compared to the tree based on the sequence of the 16S–23S rDNA Internal Transcribed Spacer (ITS) (Figure S1). Once again, good overall congruence was found between all these trees, apart from the PCC 9717 strain, which moved from SC3 to SC1 in the ITS tree. Regarding the geographical origin of the strains, it appeared that three subclades (SC2–SC4) contained strains isolated from various continents, suggesting a worldwide distribution of these strains. In contrast, SC1 only contained the two African strains, suggesting a possibly more limited distribution of this subclade.

Finally, to determine whether there was any relationship between the phylogenetic position of the strains and the size of their genome, we performed a Kruskal-Wallis analysis in which we compared the four subclades with regard to the number of CDS in the genome of each strain contained in these subclades. A statistical difference was found at a 7% level, suggesting that a putative link between the phylogenetic position of the strains and the size of their genome might have been found if a larger sample of strains in each of the four subclades had been available.

Proportion of repeated sequences

The proportion of repeats (see Materials and Methods) was carried out on the twelve M. aeruginosa genomes plus a selection of 18 cyanobacterial genomes available in our database. From this analysis, it appears that all the M. aeruginosa genomes displayed a higher proportion of repeated sequences than the other cyanobacterial genomes selected for this comparison (Fig. 4). Moreover, the slope of the regression line between the size of the genomes and the proportion of their repeats was much steeper for M. aeruginosa genomes than for the others, with NIES-843 and PCC 7806 displaying the highest proportions of repeats.

We also estimated the number of replicated CDS in the twelve Microcystis genomes by taking into account two amino-acid sequence similarity levels (70 and 90%). A highly significant relationship was found between the number of CDS they contained and the number of repeated CDS ($R^2_{Pearson} = 0.87$, p<0.0001 at 70% sequence similarity; $R^2_{Pearson} = 0.82$, p<0.0001 at 90% sequence similarity level).

Synteny analysis

The synteny values between the twelve M. aeruginosa genomes range from 67% to 86% (mean value 76±4%). In order to find out whether the synteny values estimated between the twelve genomes match with the phylogenetic relationships of the corresponding strains, we performed a non-metric MDS analysis on the matrix of these values (Fig. S2). From this figure, it appears that the strains belonging to SC1 and SC2 are clearly distinguished from those belonging to SC3 and SC4. Interestingly, the three strains belonging to SC4, which form the longest branches of the phylogenetic trees (Fig. 3), are the ones most widely scattered in the MDS scatterplot. We could conclude that the phylogenetic diversification of M. aeruginosa was accompanied by chromosome rearrangements leading to a rapid decrease in synteny.

As expected, the largest synteny group shared by the twelve M. aeruginosa genomes contains genes encoding ribosomal proteins. It is worth noting here that the second largest group comprises eleven genes involved in the transport of various nutrients, such as the gene clusters cmpABCDF and ntrABCD encoding the ABC transporter complex for bicarbonate and that for nitrate, respectively, as well as several genes (including ppsS, ppsB, ppsC) involved in the transport of phosphate.

Evaluation of horizontal gene transfer

All the Microcystis genomes contained around 11% of genes displaying more than 20% difference in their GC content compared to the mean GC content of the whole genome (Fig. 5; Table S5). This proportion is intermediate between the high percentage (>17%) of these genes found in the genomes of Prochlorococcus marinus (AS9601, MIT 9301) and Trichodesmium erythraeum (IMS101), and the lowest percentage (<3%) found in Synechococcus elongatus PCC 6301. From 106 to 247 IVOMs were found in the M. aeruginosa genomes. No relationship was found between the number of IVOMs and the size of the genome (or the number of CDSs) when we considered only the ten new genomes and the PCC 7806 strain. However, a positive correlation was found when the NIES-843 genome was included in the analysis. This correlation might be due to the fact that the NIES-843 genome is larger and contains a huge number (247) of IVOMs.

Parallel to the analysis of horizontal transfer of genetic information, we looked for genes that are strain-specific among
Table 1. General features of the ten new *Microcystis aeruginosa* genomes and of the two previously available (PCC 7806 and NIES-843).

<table>
<thead>
<tr>
<th>Strain/Culture collection reference</th>
<th>Location and date isolation</th>
<th>Genome size (Mbp)</th>
<th>SC number</th>
<th>Status</th>
<th>GC%</th>
<th>CDS number</th>
<th>Average CDS length (bp)</th>
<th>IG length (bp)</th>
<th>Coding density</th>
<th>rDNA clusters number</th>
<th>Average gene length (bp)</th>
<th>Ref</th>
</tr>
</thead>
<tbody>
<tr>
<td>PCC 7941</td>
<td>Little Rideau Lake, CAN (1954)</td>
<td>4.8</td>
<td>77</td>
<td>WGS: CAIK0000000–CAIK01000372</td>
<td>42.6</td>
<td>4669</td>
<td>839</td>
<td>227</td>
<td>82.1</td>
<td>1</td>
<td>1019</td>
<td>This work</td>
</tr>
<tr>
<td>PCC9432</td>
<td>Little Rideau Lake, CAN (1954)</td>
<td>5.0</td>
<td>132</td>
<td>WGS: CAIH0000000–CAIH01000372</td>
<td>42.5</td>
<td>4952</td>
<td>822</td>
<td>235</td>
<td>81.9</td>
<td>1</td>
<td>1000</td>
<td>This work</td>
</tr>
<tr>
<td>PCC 9443</td>
<td>Fish pond, Landjia, RCA (1994)</td>
<td>5.1</td>
<td>221</td>
<td>WGS: CAUJ0000000–CAUJ01000735</td>
<td>42.8</td>
<td>5055</td>
<td>818</td>
<td>274</td>
<td>80.6</td>
<td>1</td>
<td>1011</td>
<td>This work</td>
</tr>
<tr>
<td>PCC 9701</td>
<td>Guerlesquin dam, F (1996)</td>
<td>4.7</td>
<td>172</td>
<td>WGS: CAUO1000001–CAUO1000550</td>
<td>42.8</td>
<td>4668</td>
<td>829</td>
<td>251</td>
<td>81.8</td>
<td>1</td>
<td>1010</td>
<td>This work</td>
</tr>
<tr>
<td>PCC 9717</td>
<td>Rochereau dam, F (1996)</td>
<td>5.2</td>
<td>264</td>
<td>WGS: CAI0000000–CAI01000881</td>
<td>42.8</td>
<td>5164</td>
<td>807</td>
<td>295</td>
<td>79.7</td>
<td>1</td>
<td>1009</td>
<td>This work</td>
</tr>
<tr>
<td>PCC 9806</td>
<td>Oskosh, USA (1975)</td>
<td>4.2</td>
<td>93</td>
<td>WGS: CAIU0000000–CAIU0100236</td>
<td>43.1</td>
<td>4123</td>
<td>833</td>
<td>242</td>
<td>80.9</td>
<td>1</td>
<td>1026</td>
<td>This work</td>
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<td>5.1</td>
<td>267</td>
<td>WGS: CAIM0000000–CAIM01000236</td>
<td>42.7</td>
<td>5049</td>
<td>811</td>
<td>286</td>
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<td>141</td>
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<td>42.4</td>
<td>5044</td>
<td>811</td>
<td>241</td>
<td>81.4</td>
<td>1</td>
<td>992</td>
<td>This work</td>
</tr>
<tr>
<td>PCC 9809</td>
<td>Lake Michigan, USA 4.9 (1982)</td>
<td>4.9</td>
<td>303</td>
<td>WGS: CAIO0000000–CAIO01000789</td>
<td>42.8</td>
<td>4990</td>
<td>806</td>
<td>279</td>
<td>81.3</td>
<td>1</td>
<td>988</td>
<td>This work</td>
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<tr>
<td>T1-4</td>
<td>Bangkok, T (NA)</td>
<td>4.7</td>
<td>145</td>
<td>WGS: CAIP0100001–CAIP01000449</td>
<td>42.8</td>
<td>4610</td>
<td>830</td>
<td>251</td>
<td>81.3</td>
<td>1</td>
<td>1056</td>
<td>This work</td>
</tr>
<tr>
<td>PCC 7806</td>
<td>Baakman Reservoir, 5.3 NL (1972)</td>
<td>5.3</td>
<td>156</td>
<td>WGS: AM778843–AM778958</td>
<td>42.0</td>
<td>5496</td>
<td>775</td>
<td>215</td>
<td>81.5</td>
<td>1</td>
<td>966</td>
<td>[19]</td>
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<tr>
<td>NIES-843</td>
<td>Lake Kasumigaura, J5.8 (1972)</td>
<td>5.8</td>
<td>NC_010296</td>
<td>42.3</td>
<td>7199</td>
<td>682</td>
<td>159</td>
<td>83.0</td>
<td>2</td>
<td>812</td>
<td>[18]</td>
<td></td>
</tr>
</tbody>
</table>

Sc number: Number of scaffolds; GC%: Percentage of G/C bases in the genome; CDS number: Number of Coding DNA Sequence; IG length: Average intergenic length; Ref: Reference.

doi:10.1371/journal.pone.0070747.t001
Figure 1. Estimation of the sizes of the pangenome (A) and core genome (B) of *Microcystis aeruginosa* from the twelve *Microcystis aeruginosa* genomes (including the two previously-available genomes of PCC 7806 and NIES-843). The upper and lower edges of the boxes indicate the first quartile and third quartile, respectively, of all different input orders (1000) of the genomes. The central horizontal line indicates the sample median.
the twelve *M. aeruginosa* genomes. From 100 to 225 strain-specific genes were found in eleven of the *M. aeruginosa* genomes, and as many as 805 in NIES-843. Less than 10% of these strain-specific genes displayed over 40% identity between their deduced amino acid sequences and sequences from the Swissprot database.

Among them, we found for example, in PCC 7806, a strain-specific cluster (IPF 1564–1566) containing three genes involved in the biosynthesis of sucrose (*sppA*, *susA* and *spsA*). These three genes displayed a 40–70% amino-acid sequence identity with sequences found in various cyanobacterial genera (*e.g.*, *Nodularia*, *Nostoc*, *Cyanothece*...). As expected, we found in T1-4, the only strain among the 12 sequenced *Microcystis*, which synthesizes phycoerythrin [32], the genes required for the biosynthesis of this red phycobiliprotein and those involved in the regulation of their expression (data not shown). A 19-kb gene cluster (IPF_1564–1566) encoding for unknown proteins was identified only in the *M. aeruginosa* PCC 9808 genome. This cluster was also found with a similar physical organization and high amino acid sequence identities, in *Lyngbya majuscula* 3L (*Cyanobacteria*; Taxonomy ID: 489825) and *Herpetosiphon aurantiacus* DSM 785 (*Chloroflexi*; Taxonomy ID: 316274) (Fig. S3), and with lower amino acid sequence identities in the genomes of several proteobacteria and of two Archaea. All the genes belonging to these three clusters display a 20% deviation in their GC content, compared to the mean GC content of the genome of the strains in which they have been found.

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**Figure 2. Distribution of the core and flexible genes from all the *Microcystis* genomes in the Clusters of Orthologous Groups (COGs).** Only the COG categories containing >1% of the genes in at least one of the two core genomes, are shown in the figure. The functional classifications of the COGs are: **Cellular process and signaling**: (D) Cell cycle control, cell division, chromosome partitioning; (M) Cell wall/membrane/envelope biogenesis; (N) Cell motility; (O) Post-translational modification, protein turnover, and chaperones; (T) Signal transduction mechanisms; (U) Intracellular trafficking, secretion, and vesicular transport; (V) Defense mechanisms; (W) Extracellular structures; (Y) Nuclear structure; (Z) Cytoskeleton. **Information storage and processing**: (J) Translation, ribosomal structure and biogenesis; (K) Transcription; (L) Replication, recombination and repair. **Metabolism**: (C) Energy production and conversion; (E) Amino acid transport and metabolism; (F) Nucleotide transport and metabolism; (G) Carbohydrate transport and metabolism; (H) Coenzyme transport and metabolism; (I) Lipid transport and metabolism; (P) Inorganic ion transport and metabolism; (Q) Secondary metabolites biosynthesis, transport, and catabolism. **Poorly characterized**: (R) General function prediction only; (S) Function unknown.

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Secondary metabolites

Eleven gene clusters encoding non-ribosomal peptide synthetase (NRPS) and/or polyketide synthase (PKS), and two ribosomal ones predicted to be involved in the biosynthesis of secondary metabolites, were found among the *Microcystis* genomes (Table 2). Each genome contained between two and nine such gene clusters accounting for 1.0 to 3.4% of the total genome size. Seven of these clusters encode enzymes for the biosynthesis of known metabolites (microcystins, aeruginosins, cyanopeptolins, microginins, anabaenopeptins, cyanobactins and microviridins), whereas the six remaining clusters encode enzymes for the biosynthesis of so-far unidentified products (Fig. 6).

Four of these gene clusters (cyanopeptolins, aeruginosins, microcystin and microviridins) were found in most of the genomes (Table 2), and their deduced amino acid sequences displayed high percentages of amino-acid sequence identity (AASI, >90%). However, some variability was found in the aer gene cluster, both in its composition (presence/absence of aerJ and aerM for instance) and in the length of the gene sequences (1578 to 4287 bp long for aerA in PCC 9432 and PCC 9807). With regard to the mdn gene cluster, the MdhB-E proteins were highly conserved (AASI, >90%), whereas MdhA, which is involved in precursor synthesis, was less conserved (>52%). Although some mdn genes were lost from the PCC 9806 genome, the presence of mdnB and mdnD in this genome indicated that the microviridin gene cluster was originally present in all the *Microcystis* genomes examined.

All the other gene clusters potentially involved in the synthesis of secondary metabolites were found in only one to four genomes. For example, the proteins involved in the biosynthesis of anabaenopeptins were extremely well conserved (AASI, >97%) in PCC 9432 and PCC 9701. On the other hand, the proteins involved in the biosynthesis of microginin and of the cyanobactins (such as microcyclamides) appeared to be less conserved (AASI, 85% and >82%, respectively). Similarly, the cyanobactin gene cluster of PCC 9432 displayed considerable differences from that present in PCC 7806.

In addition to these already well-known gene clusters, six orphan gene clusters were also retrieved (Table 2, Fig. 6). The extremities of these gene clusters have been defined by using the available data from the first genome in which they have been found (e.g. PKSImod/PKSI, PKSI-I iterative in Frangeul et al. [19]). For example, the high synteny and deduced amino acid sequence identity (cluster of 12 adjacent genes with AASI, >99.2%) found between the PKSImodular/PKSIII gene clusters in four *M.*
aeruginosa genomes revealed that this cluster contained at its 3’ end, some additional genes to the ones previously described for this cluster [19] (Fig. 6).

Three new putative NRPS and/or PKS gene clusters (MIC1, MIC2 and MIC3) were retrieved from at least three genomes with high AASI (>92%) and may be involved in the production of so-far unknown compounds. MIC1 is the most promising gene cluster, with adjacent NRPS and PKS genes and RND superfamily efflux transporters. MIC2 and MIC3 were organized as individual thiotemplate modular systems surrounded by putative tailoring enzymes.

Finally, a Correspondence Analysis performed on the distribution of all these gene clusters among the twelve M. aeruginosa genomes revealed that three (SC1, SC2 and SC3) of the four subclades defined by our phylogenetic approach cannot be distinguished by their content in these genes (Fig. S4). On the other hand, the three genomes belonging to SC4 are clearly differentiated from the other genomes by the absence of mcy and aer gene clusters and more generally by their low content of genes involved in the biosynthesis of secondary metabolites.

**Discussion**

Despite the multiple sequencing strategies used in this study and the very high sequence coverage, the final assembly process was impeded by the very high proportions of repeated sequences present in the ten new M. aeruginosa genomes. However, several observations indicate that a comparative genomic analysis could be undertaken with good confidence. First, the percentage of 454 reads (around 97%) used for assembly was in the same range than that obtained for finished genomes. Second, no gene was lacking in the Minimal Gene Set of the ten new genomes, when compared with that of the M. aeruginosa NIES-843 complete genome. Third, an almost asymptotic curve for the core genome was reached, a result that could not be obtained if numerous genes had been missing in these genomes.

The adaptive capacities of M. aeruginosa, which may explain its worldwide distribution and its ability to proliferate and to dominate the phytoplankton communities in eutrophic freshwater ecosystems, seem to rely on a particular genome evolutionary strategy. This strategy combines a large genome plasticity, characterized by a high number of repeated sequences, numerous rearrangements and an ability to include new adaptive genes by horizontal gene transfer.

Compared to other cyanobacterial genomes the M. aeruginosa genomes display both high proportions of repeated sequences and wide variations in their proportions depending on the strains. The proportions of these repeats appear to be lower in the ten new M. aeruginosa genomes than in the two previously-known ones obtained by a Sanger-sequencing approach. The Newbler assembler used for the assembly of the ten new genomes is known to span a lot of repeats from the contigs [33], and so it is likely that we have underestimated the proportions of repeats in the ten new genomes. As previously described by Larsson *et al.* [34], who...
compared the genomes of different cyanobacterial genera, we also found a positive correlation between the genome size and the number of duplicated genes. It has been hypothesized that a combination of both gene duplication and transfer of orthologous alien genes, without replacement of the native genes, led to an increase in genetic redundancy that contributes to the robustness of biological systems, i.e. their ability to continue to function despite external and internal perturbations (e.g. [35]). A contrario, a loss of genetic redundancy in bacteria displaying a reductive genome evolution, as observed in numerous pathogens [36] and in oceanic picocyanobacteria of the genus Prochlorococcus [15], would result in weak selection for robustness, reflecting the fact that these microorganisms live in stable environments. Thus, the unusually high genetic redundancy in the *M. aeruginosa* genomes might be regarded as an evolutionary strategy that allows this cyanobacterial species to occupy changing environments, such as freshwater ecosystems.

Another peculiar trait of *M. aeruginosa* concerns the low synteny values found for the twelve genomes analyzed in the present study. These low synteny values contrast with the very high 16S rRNA sequence similarities (>99.5%) found between *M. aeruginosa* genomes, and indicate a rapid evolution of the gene organization that probably permits to generate new combinations of genes allowing different adaptive capacities to emerge.

The third feature of the *M. aeruginosa* genomes is that *M. aeruginosa*, like Prochlorococcus [14,37], displays an open pangenome that allows it to acquire new genes. Among them, it was very interesting for example to find the gene cluster involved in the biosynthesis of sucrose in *M. aeruginosa* PCC 7806 alone, because sucrose is known to be involved in the salt acclimation mechanism [38]. Indeed, Guljamow et al. [39] had identified actin and profilin genes in PCC 7806 genome, which could be involved in the adaptation to high osmotic stress. Thus, our finding reinforces the hypothesis that PCC 7806 cells may have acquired by HGT two gene clusters that have allowed them to survive and develop in their original habitat (Braakman water reservoir, The Netherlands), which is known to have undergone successive changes in the salt concentration during its history [39]. Another interesting finding was the presence in *M. aeruginosa* PCC 9808 of a large gene cluster (19 kb, IPF_1564–1566) potentially resulting from HGT that illustrates the ability of *M. aeruginosa* to integrate large DNA molecules into its chromosome.

The plasticity of the *M. aeruginosa* genome seems to be linked to the ability of members of this species to bloom in a wide range of ecosystems worldwide. However, such a strategy results in disorder in the organization of the genome, and might also be costly for cell functioning. For example, it has been shown that maintaining redundant genes is costly when these genes belong to the core...
Figure 6. Schematic representation of all the secondary metabolite gene clusters found in the twelve *Microcystis aeruginosa* genomes (including the two previously-available genomes of PCC 7806 and NIES-843). For each biosynthesis cluster, the sketch corresponds to the gene cluster present in the reference strain genome, its size in kb and the amino acid sequence identity estimated for the orthologous region in the other *Microcystis* genomes. The reference strain genome corresponds to the one indicated in Table 2.

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Table 2. Distribution and amino-acid sequence identity of 13 gene clusters involved in the biosynthesis of secondary metabolites in the twelve *Microcystis* genomes (including the two previously available, PCC 7806 and NIES-843).

<table>
<thead>
<tr>
<th>Strain</th>
<th>% of genome</th>
<th>Nb clust/strain</th>
<th>mcy seq ident (Detection)</th>
<th>aer seq ident (Detection)</th>
<th>mcn seq ident (Detection)</th>
<th>mic seq ident (Detection)</th>
<th>apt seq ident</th>
<th>mca seq ident</th>
<th>mdn seq ident (Detection)</th>
<th>Ref.</th>
<th>Aa seq ident for mdnB-E</th>
<th>Aa seq ident to AB279593</th>
<th>Aa seq ident</th>
<th>Aa seq ident</th>
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<tr>
<td>PCC 7941</td>
<td>3.3</td>
<td>6</td>
<td>98.3 (+)</td>
<td>94.9 (−)</td>
<td>95.3 (+)</td>
<td>Ref.</td>
<td>-</td>
<td>-</td>
<td>98.2</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>PCC 9432</td>
<td>3.4</td>
<td>9</td>
<td>- (−)</td>
<td>94.8 (+)</td>
<td>91.2 (+)</td>
<td>89.9*</td>
<td>Ref.</td>
<td>81.7*</td>
<td>98.2</td>
<td>99.2</td>
<td>partial£</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>PCC 9443</td>
<td>2.3</td>
<td>4</td>
<td>96.9 (na)</td>
<td>95.5 (+)</td>
<td>mcnG-F£ (-)</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>90.0</td>
<td>96.1</td>
<td>-</td>
<td>-</td>
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<td>-</td>
</tr>
<tr>
<td>PCC 9701</td>
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<td>4</td>
<td>- (na)</td>
<td>- (na)</td>
<td>90.7* (+)</td>
<td>85.5</td>
<td>97.3</td>
<td>-</td>
<td>98.5</td>
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<tr>
<td>PCC 9717</td>
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<td>6</td>
<td>mcyA-C£ (na)</td>
<td>93.4 (na)</td>
<td>91.3* (+)</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>98.1</td>
<td>-</td>
<td>-</td>
<td>98.2</td>
<td>95.5*</td>
<td>98.6*</td>
<td></td>
</tr>
<tr>
<td>PCC 9806</td>
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<td>2</td>
<td>- (na)</td>
<td>- (na)</td>
<td>- (na)</td>
<td>-</td>
<td>-</td>
<td>91.5</td>
<td>mdrB£</td>
<td>-</td>
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<td>PCC 9807</td>
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<td>5</td>
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<td>90.2 (+)</td>
<td>96.1(*)</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>96.2</td>
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<td>-</td>
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<td>-</td>
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<td>6</td>
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<td>93.2 (+)</td>
<td>92.3 (−)</td>
<td>-</td>
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<td>98.1</td>
<td>-</td>
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<td>93.5 (+)</td>
<td>92.0* (+)</td>
<td>-</td>
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<td>98.6</td>
<td>99.3</td>
<td>-</td>
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<td>T1-4</td>
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<td>3</td>
<td>- (na)</td>
<td>- (na)</td>
<td>- (na)</td>
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<td>-</td>
<td>-</td>
<td>97.1</td>
<td>-</td>
<td>94.9</td>
<td>96.2</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>PCC 7806</td>
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<td>7</td>
<td>Ref. (+)</td>
<td>Ref. (+)</td>
<td>Ref. (+)</td>
<td>Ref.</td>
<td>Ref.</td>
<td>Ref.</td>
<td>96.6</td>
<td>Ref.</td>
<td>Ref.</td>
<td>Ref.</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>NIES-843</td>
<td>2.5</td>
<td>5</td>
<td>96.6 (+)</td>
<td>93.8 (na)</td>
<td>91.6 (na)</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>Ref.</td>
<td>Ref.</td>
<td>Ref.</td>
<td>Ref.</td>
<td>-</td>
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</tr>
</tbody>
</table>

| No of strains containing the complete gene cluster | 9 | 9 | 3 | 2 | 4 | 11 | 4 | 2 | 1 | 4 | 3 | 3 |

mcy: Microcystins; aer: Aeruginosins; mcn: Cyanopeptolins; mic: Microginins; apt: Anabaenopeptins; mca: Cyanobactins; mdn: Microviridins; PKSI iterat: PKSI iterative; % of genome: Proportion of genome; Nb clusters/strain: Number of clusters per strain. Aa seq ident: Amino-acid sequence identity; Ref.: Reference strain used for the estimation of the sequence identity. £: incomplete gene cluster; *: on adjacent contigs highly fragmented genome or region; +: metabolite detected in previous studies ([58]; Welker, unpublished); £: probably encodes for another cyanobactin; (+): positive detection of the metabolite in the strain; (−): negative detection of the metabolite in the strain; na: not analyzed. doi:10.1371/journal.pone.0070747.t002
Among the 13 gene clusters involved in the biosynthesis of secondary metabolites, microcystins, aeruginosins, cyanopeptolins and microviridins were those most widely distributed in the twelve *Microcystis* genomes, but it appeared that one subclade (SC4) was clearly distinguishable from the others. These three strains displayed the longest branches in the phylogenetic tree and originated from different continents, suggesting that these widely distributed NRPS/PKS gene clusters were lost early in the evolutionary radiation of these strains. The common distribution of the microcystin, aeruginosin and cyanopeptolin metabolites in three other subclades compared to the more sporadic distribution of the other secondary metabolites, suggest that putative interactions lead to their joint conservation. The observation that their gene expression profiles during a day/night cycle are similar [56], is consistent with the above hypothesis.

In conclusion, it seems interesting to compare the main features of the *M. aeruginosa* genomes with those of *P. marinus*, because members of these two cyanobacterial species are characterized by having wide geographical distributions, and play major roles in their respective environments (freshwater versus oceanic ecosystems). From this comparison it appears that members of both species display very different evolutionary strategies even if some common features can be found in their genome organization. For example, among the twelve genomes contain almost the same proportions of core and flexible genes, even if the mean size of the *M. aeruginosa* genomes exceeds more than two fold that of the *P. marinus* genomes. In the same way, both species display an open pangenome that permits them to colonize various ecological niches.

But in contrast with *Prochlorococcus*, *M. aeruginosa* does not seem to be organized in ecotypes, and its large genome contains a large number of repeat sequences and a high proportion of transposases. These characteristics might permit rapid variation in gene content and the occurrence of new gene combinations allowing *M. aeruginosa* populations to cope with various environmental conditions encountered by this species. Moreover, the *M. aeruginosa* genomes harbor wide genetic diversity with ribosomal and non-ribosomal gene clusters dedicated to synthesis of these bioactive compounds whereas the reduced genomes of *P. marinus* are nearly devoid of all these secondary metabolite biosynthesis gene clusters, even if it has been described that some *Prochlorococcus* strains are able to produce various lantipeptides by using a single biosynthetic enzyme gene [57]. Finally, it is interesting to consider the sharply contrasting evolutionary strategies adopted by *M. aeruginosa* and *P. marinus*, and their ability to dominate their respective habitats.

Freshwater ecosystems are very small in comparison to oceanic areas, and consequently are more likely to display rapid fluctuations in their physico-chemical and biological characteristics. Knowing that the differentiation of ecotypes is linked to the occupation of a stable niche, *M. aeruginosa* has developed another winning genomic strategy based on the high plasticity of its genome, to enable it to cope with and colonize unstable freshwater ecosystems efficiently.

**Supporting Information**

**Table S1** Overall technical characteristics of the ten new *Microcystis aeruginosa* genomes.

( DOCX)

**Table S2** Distribution of the tRNA genes in the twelve *Microcystis aeruginosa* genomes (including the two previously-available genomes of PCC 7806 and NIES-843).

( DOCX)
Table S3  Distribution of the 205 genes from the Minimal Gene Set [30] in the twelve Microcystis aeruginosa genomes (including the two previously-available genomes of PCC 7806 and NIES-843).

Table S4  Automatic annotation and COG assignation of the Microcystis aeruginosa core genes.

Table S5  List of the genes, for the twelve Microcystis genomes, displaying more than 20% difference in their GC content compared to the mean GC content of the whole genome.

Figure S1  Phylogenetic relationships (Neighbor-joining method) between the twelve Microcystis aeruginosa genomes (including the two previously-available genomes of PCC 7806 and NIES-843) based on the alignment of the 16S–23S rDNA Internal Transcribed Spacer.

Figure S2  Non-metric MDS analysis performed on the matrix of synteny values estimated between the twelve Microcystis aeruginosa genomes (including the two previously-available genomes of PCC 7806 and NIES-843). SC1, 2, 3 & 4: Sub-clades defined in the phylogenetic tree based on the core genome (see Fig. 3).

References


