Different behaviour between autotrophic and heterotrophic *Galdieria sulphuraria* (Rhodophyta) cells to nitrogen starvation and restoration. Impact on pigment and free amino acid contents

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**Abstract**

The unicellular red alga *Galdieria sulphuraria* is a polyextremophilic organism with a metabolic flexibility to grow autotrophically or heterotrophically. *Galdieria* can also produce and accumulate biotechnologically attractive products such as pigments (phycocyanin) and proteins. In this research we studied the effects of nitrogen starvation and its subsequent restoration on pigment and free amino acid contents both in photoautotrophic and heterotrophic cells. Following the nitrogen starvation, the levels of the primary photosynthetic pigments decreased both in autotrophic and heterotrophic cells, except for the chlorophyll *a* marginally diminished in heterotrophic cells. Ammonium supply to *G. sulphuraria* N starved cells caused a significant increase of total chlorophylls both in autotrophic and heterotrophic cells. It was observed how such increase was more rapid and marked in heterotrophic cells than in the autotrophic ones. Under N starvation, phycocyanin contents decreased in both autotrophic and in heterotrophic cells; however, after a time-lapse of 24 hours, they resulted significantly higher in heterotrophic cells. In *Galdieria sulphuraria*, like in other microalgae, free amino acid contents were profoundly dependent on nitrogen status of the cells but heterotrophic cells maintained much higher levels, especially of glutamate, respect to autotrophic ones. In general, cells grown in the presence and absence of light showed different responses toward N availability; in particular heterotrophic cells seemed to respond quicker to the ammonium restoration compared to autotrophic ones.

**Introduction**

Among abiotic factors capable of influencing the algal growth, the availability of nutrients is one of the most important, along with light and temperature. Microalgae, as well as plants, need macro- and micronutrients for their growth. Macronutrients, such as nitrogen (N), are required in larger quantities; in fact, they represent structural components of cells and are important for osmoregulation. Nutrient deficiency, in particular nitrogen, affects many aspects of plants and microalgae physiology. General symptoms of N deficiency, like the decreasing growth rate, the decline in cell protein and in pigment contents, have been all well described in microalgae. Generally, microalgae and cyanobacteria maintain a restricted ability to preserve the photosynthetic functions during N starvation. Also, in microalgae, the effects due to the lack of a nutrient can be rapid (hours) and mediated by an oxidative imbalance.

The unicellular red alga *Galdieria sulphuraria* is a member of the Cyanidiales, polyextremophilic microalgal thriving in highly acidic environments (pH 0.05-5.00) with elevated temperatures up to 56 °C, where *Galdieria* is the predominant genus. The horizontally acquired genes of *G. sulphuraria* with the ability to maintain neutral cytosolic pHs despite the highly acid environment, as well as salinity and metal resistance. Besides, many metabolite transporters are encoded in the genome of *G. sulphuraria*, explain the unusual ability to survive in such extreme habitats. For example, the horizontal acquisition of several bacterial ion pumps is recognized to provide *G. sulphuraria* with the ability to maintain neutral cytosolic pHs despite the highly acidic environment.

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attention of many researchers for its excellent properties and its likely use in many biotechnological applications. *Galdieria sulphuraria* C-PC is not only very resistant to high temperature but can also be used as a food additive colouring agent, and as a natural antioxidant ingredient and for others biotechnological purposes.

The growth and the biochemical composition of microalgae are extremely sensitive to alterations in physical and chemical parameters of the culture medium. For example, the photosynthetic pigment levels vary depending on the cultivation conditions such as light regime, temperature, inorganic carbon content, medium pH or stress. In fact, by manipulating culture conditions, it is possible to get higher or lower concentrations of a certain pigment. The N natural resources for autotrophic organisms are ammonium and/or nitrate. The absorbed nitrate is firstly reduced to ammonium and then assimilated into glutamine through the glutamine synthetase enzymes. Once assimilated as glutamine, nitrogen is transferred to 2-oxoglutarate to form glutamate through the glutamate synthase, and further utilized to form various nitrogen organic compounds. Although the matter is still debated, glutamate dehydrogenase could also be used in
the nitrogen assimilation.\textsuperscript{27} \textit{Galdieria sulphuraria}, as all members of \textit{Cyanidiales}, can assimilate nitrogen from the culture medium as nitrate or ammonium salts. Therefore, the main nitrogen source in acidic hot springs is the ammonium and nitrate assimilation rarely occur in these environments.\textsuperscript{28} The effect of nitrogen starvation on photosynthetic pigments was previously considered in green algae\textsuperscript{29-32} and in cyanobacteria.\textsuperscript{3} In \textit{Cyanidiales}, only the effects of a long-term N starvation (10 days) in autotrophic cells had been taken into account by.\textsuperscript{14} The aim of this paper is to investigate and clarify the short-term effects of N starvation (24 h) and its subsequent resupply (24 h) on pigment and free amino acid contents in both autotrophic and heterotrophic cells of \textit{G. sulphuraria}.

### Materials and Methods

#### Algal Strains and Growth Conditions

Experiments were performed with pure cultures of the red algae \textit{Galdieria sulphuraria} (strain 011) from the ACUF collection of the Department of Biology of the University of Federico II, Naples, Italy (http://www.biologiavegetale.unina.it/acuf.html). Cells were grown in batches containing 1 L of Allen medium\textsuperscript{33} with 10 mM (NH$_4$)$_2$SO$_4$ as nitrogen source. They were placed in a thermostatic chamber (Angelantoni CH 770) under continuous irradiance 90 μmol m$^{-2}$ s$^{-1}$ photons at 38 ± 1°C. The pH was set at 1.5 and controlled daily. Carbon dioxide was supplemented by filtered air blowing into the medium. Furthermore, each batch containing the heterotrophic cultures was wrapped with aluminum foil to ensure that the cultivation took place in the dark, and organic carbon was supplemented with 2% (w/v) glycerol.

In the experiments of N starvation, the cells were harvested during the logarithmic phase of growth (culture OD$_{600}$ ~ 1.0) by low speed centrifugation at 4000 × g for 10 min and washed twice with N-free Allen’s medium where (NH$_4$)$_2$SO$_4$ was omitted. Then, the supernatant was removed, and the algal pellets were re-suspended in N-free Allen’s medium and cultured up to 24 h. The effects of the ammonium addition to the nutrient solution of N starved cells for 24 hours were analysed supplying (NH$_4$)$_2$SO$_4$ (4000 × g for 15 min). The packed cells were treated with 1 mL of cold 80% ethanol and vortexed. The amino acids were extracted for 10 min at 4°C and finally clarified by centrifugation. The supernatant was filtered using Waters Sep-Pak Cartridges C18 light (Milan, Italy), and utilized for amino acid analysis as previously described.\textsuperscript{34} Quantification of glutamic acid (Glu), asparagine (Asn) and glutamine (Gln) were made against a relative calibration curve and expressed as μmol mL$^{-1}$ PCV.

### Extraction and determination of amino acids

Samples of the cell suspensions (10 mL) were collected at given times by centrifugation.28 The effect of nitrogen starvation on photosynthetic pigments was previously considered in green algae29-32 and in cyanobacteria.3 In \textit{Cyanidiales}, only the effects of a long-term N starvation (10 days) in autotrophic cells had been taken into account by.14 The aim of this paper is to investigate and clarify the short-term effects of N starvation (24 h) and its subsequent resupply (24 h) on pigment and free amino acid contents in both autotrophic and heterotrophic cells of \textit{G. sulphuraria}.

### Spectroscopic estimation of pigments in the crude extracts

Chlorophyll (Chl) (a and total) were extracted with N, N-dimethylformamide. The Chl (a and total) contents were estimated spectrophotometrically at 647 and 665 nm according to Carfagna et al.\textsuperscript{35} The C-PC contents were estimated as previously described.\textsuperscript{10}

### Packed cell volume (PCV) determination

Packed Cell Volume (PCV) was measured by centrifuging a known aliquot of cell suspension in a hematocrit tube at 4000 g for 5 min as previously reported.\textsuperscript{34} The packed cell volume (PCV) was measured by centrifuging a known aliquot of cell suspension in a hematocrit tube at 4000 g for 5 min as previously reported.\textsuperscript{34}

### Statistical analyses

Experimental data analysis was carried out using the Sigmplot 14 software. Data of the mean ± standard deviation of three independent experiments were also presented. The statistical analysis was performed by one-way analysis of variance (ANOVA) with a Tukey post ad hoc test to determine differences between autotrophic and heterotrophic conditions.

### Results and Discussion

After carbon, nitrogen is the most essential nutrient for plant and algae since it contributes directly to biomass production. The availability of N affects the photosynthesis activity in a way that low quantities of nitrogen compromises the structure of the photosystems in terms of protein and photosynthetic pigment contents. In this study, we consider and compare the short-term effect of N starvation and restoration in \textit{G. sulphuraria} cells both in autotrophic and in heterotrophic growth condition. \textit{Galdieria} cells, grown in a complete Allen medium and kept in the light (autotrophic control cells) showed similar contents of Chl $a$ (3.9 ±0.2 μg μL$^{-1}$ PCV) and total Chl (5.6 ± 0.4 μg μL$^{-1}$ PCV) if related to heterotrophic control cells (Chl $a$, 2.7 ±0.3 μg μL$^{-1}$ PCV; total Chl, 4.9 ± 0.3 μg μL$^{-1}$ PCV) (Figure 1A-B). Most \textit{Galdieria} strains lose
their pigmentation when they are heterotrophically grown.\textsuperscript{13,36} Unlike the others, \textit{Galdieria sulphuraria} (strain ACUF 011) does not bleach under heterotrophy since Chls levels are not considerably different between auto- and heterotrophic cells. Anyway, total Chl decreased in autotrophic cells under N starvation while it remained almost unchanged in heterotrophic cells (Figure 1B) with Chl \textit{a} content strongly decreased already after 24 hours of deprivation (Figure 1A). Ammonium supply, following the 24h of N starvation, caused a significant increase of total Chl both in autotrophic and heterotrophic cells (Figure 1B). Therefore, the increase in total Chl was observed to be more rapid and marked in heterotrophic cells. In fact, after only two hours from the ammonium supply, Chls were 9.98 ± 0.5 µg µL\textsuperscript{-1} PCV, values significantly higher respect to those of autotrophic cells (Figure 1B). The amount of total Chl was similar in both auto- and heterotrophic cells after six hours from the N supply and dropped in the following hours (Figure 1B).

Nitrogen assimilation and photosynthetic pigments synthesis are closely linked. The loss of pigments (i.e. chlorophylls, phycobiliproteins) caused by N starvation is considered, together with cessation of cell division, a general response that occurs in all photoautotrophic organisms.\textsuperscript{7,31} In autotrophic cells, the decrease in Chls level is very harmful as it reduces the ability to capture light and to perform an efficient photosynthesis. Chlorophylls are N-containing-macromolecule and a reduction of their ex-novo synthesis lead to a decrease in the nitrogen demand. Typically, in microalgae, the diminution in Chl content leads to a dramatic change in colour from usual blue-green to yellow-green. Such phenomenon is referred to as bleaching or chlorosis.

As previously reported by Perez-Garcia \textit{et al.},\textsuperscript{37} photosynthetic pigments in the light-harvesting complex in \textit{G. sulphuraria} under heterotrophic cultivation were slightly lower than those cultivated with photoautotrophic cultivation. These lower contents have been attributed to the reduction of their synthesis since the energy, in heterotrophic cells, derives from oxidation of the organic substrate and not from photosynthesis.

Our results indicate that generally the pigments decrease under N deprivation both in autotrophic and heterotrophic cells, but with a different time course and to a different extent. Chl \textit{a} was rapidly affected in autotrophic cells while marginally decreased in heterotrophic cells in the first 6 hours of starvation. This result suggested that during the N deprivation the synthesis of the components of the antenna system was preferred over the photosystem reaction centres. It is reasonable to think that these pigments have not only the role of an antenna, and therefore linked to the capture of light for photosynthesis, but they are also used as N reserve. Also, total Chl, that in the \textit{Cyanothecales} include chlorophyll \textit{a} and \textit{d},\textsuperscript{38} was more affected by N starvation in autotrophic cells respect to heterotrophic ones.

In this study also the change in C-PC levels were considered. After 6h of N starvation, C-PC contents decreased significantly (P≤0.05) in autotrophic (from 10.7±0.5 to 3.8±0.2 µg µL\textsuperscript{-1} PCV) and in heterotrophic cells (from 6.9±0.07 to 1.4±0.2 µg µL\textsuperscript{-1} PCV). After 24 hours, C-PC continued decreasing (2.7±0.4 µg µL\textsuperscript{-1} PCV) while resulted significantly increased in heterotrophic cells reaching the value of 12.3±1.6 µg µL\textsuperscript{-1} PCV. Ammonium addition provoked a sudden increase in C-PC in both cell types. After only two hours from the addition of ammonium, it was noted an increase in C-PC, reaching the value of 8.3±1.4 µg µL\textsuperscript{-1} PCV in autotrophic cells and 22.4±1.0 µg µL\textsuperscript{-1} PCV in heterotrophic ones (Figure 2). Therefore, in the following four hours of N restoration, C-PC continued to increase in the autotrophic cells, stopping in the heterotrophic ones. In the following hours up to 24 hours from the ammonium addition, the concentration of C-PC decreased in both cells, particularly under heterotrophic condition (Figure 2).

Definitely, N-deprivation also reduced the content of C-PC for all 24 hours in autotrophic cells but only in the first 6 hours in heterotrophic cells. Indeed, in the latter, there was a sharp increase in concentration at the end of the starvation period. The results seem to indicate that N starvation most affected autotrophic cells, concerning Chls and C-PC, compared to heterotrophic cells. These observations suggested that cells used these N-containing pigments when the intracellular level of nitrogen depleted. Perhaps, these nitrogenous compounds are converted into essential N-containing cell materials such as proteins, nucleic acid, and cell wall materials necessary for the cell growth or the synthesis of polypeptides required for acclimation to the N starvation.\textsuperscript{14,39} Moreover, C-PC could act as a nitrogen store as previously reported by Sloth \textit{et al.}\textsuperscript{39} while its degradation would be rather for minimising the absorption of excess excitation energy under stress conditions as in cyanobacteria.\textsuperscript{40} In \textit{Synechococcus}, a C-PC decrease occurs within 8 hours of N starvation.\textsuperscript{3} Furthermore, the N deprivation limits Chls and C-PC synthesis since protoporphyrin IX, their common precursor, is a product of ammonium assimilation.\textsuperscript{21}

![Figure 2. Phycocyanin content (µg µL\textsuperscript{-1} PCV) as a function of NH\textsubscript{4}\textsuperscript{+} availability (N) in \textit{Galdieria sulphuraria} cells cultured in autotrophy, and in heterotrophy. On the left side, it is indicated the data referring to the nitrogen starvation (-N), while on the right hand side, the data after nitrogen addition (+N), indicated by the dotted line. Error bars represent standard deviation of the mean (n = 3). The same letters above each bar, within the same graph, indicate non-significant differences (P ≤ 0.05, ANOVA, Tukey’s multiple comparison).](image-url)
Sinetova et al.14 showed relevant changes occurring in the chloroplast ultrastructure in Galdieria sulphuraria photoautotrophic cells under N deficiency. The chloroplast was reduced, and storage compounds accumulated. The occurrence of storage compounds in the chloroplast could explain the delayed start of nitrogen assimilation, with the addition of ammonium, which occurs in autotrophic cells of G. sulphuraria.

Free amino acid contents in the algal cell is an excellent index of the ability to assimilate the nitrogen just added to the culture.29 Typically, in N starved algal cells the content of free amino acids is reduced due to the slowdown of their synthesis. In addition, N-deprived and photosynthesizing algal cells are always very ready to assimilate ammonium as early as the first two hours after N addition. In this study, Gln and amide levels were different in autotrophic and heterotrophic cells, both under N starvation and after ammonium restoration (Table 1). The concentration of free Gln resulted much higher in the heterotrophic cells than in the autotrophic ones. Under N deprivation, Gln did not change in both auto- and heterotrophic cells, but its level was about twice as high in heterotrophic as compared to autotrophic cells. Furthermore, after 2 hours from ammonium supply, Gln decreased in autotrophic cells (from 7.10 ± 0.67 to 5.65 ± 0.15 µmol mL⁻¹ PCV), while it was about four times higher in the heterotrophic cells than in the autotrophic cells. In the following few hours, Gln contents restored in autotrophic cells while decreased in heterotrophic ones (Table 1).

The concentration of free glutamine (Gln) resulted similar in autotrophic and heterotrophic cells (Table 1). Therefore, after 6 hours of N starvation, Gln increased in both autotrophic (from 0.74 ± 0.13 to 1.93 ± 0.19 µmol mL⁻¹ PCV) and in heterotrophic cells (from 0.67 ± 0.15 to 1.80 ± 0.38 µmol mL⁻¹ PCV). After 24 hours, Gln resulted almost unchanged at 1.98 ± 0.56 µmol mL⁻¹ PCV in autotrophic cells and decreased to 0.70 ± 0.20 µmol mL⁻¹ PCV in the heterotrophic ones. Nitrogen re-supply increased the Gln content in both autotrophic and heterotrophic cells. After 2 hours of N starvation, Gln doubled (4.48 ± 0.89 µmol mL⁻¹ PCV) in both cell types (1.62 ± 0.55 µmol mL⁻¹ PCV) (Tab.1). At the end of the experiments, the Gln contents were similar in both conditions (about 7.0 µmol mL⁻¹ PCV).

N starvation for 24 hours affected Asn content, which decreased in autotrophic cells (from 2.95 ± 0.73 to 1.59 ± 0.73 µmol mL⁻¹ PCV) and increased in heterotrophic ones (from 1.16 ± 0.22 to 3.31 ± 0.50 µmol mL⁻¹ PCV). However, following the addition of ammonium, asparagine increased significantly in autotrophic cells after 2 hours, while slightly decreased in both cell types in the following hours.

In our experiments, heterotrophic cells seem to respond quickly to the ammonium addition when compared to autotrophic ones. In G. sulphuraria N starved for 24h, the Gln levels are significantly higher in autotrophic cells than in heterotrophic ones. The high content of Gln in autotrophic cells could exert an inhibiting effect of ammonium uptake as previously reported.30 High-affinity ammonium transport and AMT1.1 expression appear to be subject to down-regulation by glutamine.41 It is reasonable to suppose that the high content of Gln in autotrophic cells may exert a temporary inhibition of ammonium uptake, explaining the delayed effect (2h) of ammonium supply on changing Chl contents. Indeed, in heterotrophic cells, where Gln levels were significantly lower than autotrophic cells, the effects of ammonium addition on metabolite changing took place in a short time.

Among the intracellular free amino acids, Gln is the one that change less marginally, even when cells were exposed to N deprivation.42 In fact, Gln forms a steady cellular pool from which to draw for the synthesis of other amino acids and N compounds. Its decrease, immediately after the addition of ammonium to autotrophic cells, could indicate the activation of glutamine synthetase enzyme, which uses glutamate as a substrate. After that time, glutamate concentration was promptly restored.

In this regard, it is very interesting to note that cells in heterotrophy had much higher levels of Gln than autotrophic cells. In general, high intracellular levels of Gln may indicate poor ability to assimilate ammonium in the form of Gln. Indeed, heterotrophic cells had a lower total N content than autotrophic cells, lacking photosystems and proteins related to the photosynthetic activity.17 In addition, Gln is a constituent of glutathione together with cysteine and glycine. Cells from heterotrophic cultures of Galdieria sulphuraria and Galdieria phlegrea exhibited high levels of glutathione content;16,17 thus, high levels of glutamate could certainly be related to high levels of glutathione found in heterotrophic cells. Glutamate in heterotrophic cells remained high and unchanged after the ammonium supply (2h). However, it significantly decreased after 6 hours from the N addition. This result could be explained by the activation of the enzyme glutamate dehydrogenase to reform ammonia from α-ketoglutarate or the Krebs cycle, a prominent metabolic pathway in cells under heterotrophic growth condition.37 Another alternative is that the enzyme glutamate dehydrogenase is involved in detecting the redox status of the cell and as such, it can represent a stress monitoring protein.36 It is captivating to note that the asparagine present in cells in heterotrophy condition, to some extent, increased during the N deprivation. A future study will be conducted to evaluate the activity of phosphoenolpyruvate carboxylase enzyme in these cells, which usually supplies oxoacetate to the Krebs cycle to support the aspartate (Asn precursor) and amino acid synthesis.

Table 1. Changes of free amino acid concentrations in Galdieria sulphuraria, growing in autotrophy and heterotrophy, either under N starvation or after ammonium addition. The values reported are the average of three separate experiments±Standard Error (SE).

| Growth condition | Amino Acid | 0 Time (h) –N | 6 | 24 | Time (h) +N | 2 | 6 | 24 |
|------------------|------------|---------------|---|----|---------------|---|---|----|
| Autotrophy       | Gln        | 7.10±0.67ab   | 9.21±0.69ab | 5.65±0.15ab | 11.39±2.01ab | 7.90±1.14ab | 11.39±2.01ab | 7.90±1.14ab |
|                  | Asn        | 2.95±0.73ad   | 0.84±0.05ab | 1.50±0.73ab | 5.73±0.38ac  | 3.41±0.47ac  | 1.20±0.23ac  | 5.73±0.38ac  |
|                  | Gln        | 0.74±0.13ab   | 1.90±1.19ab | 1.98±0.56abc | 4.84±0.89ac  | 1.42±0.34bc  | 7.79±0.49Ad  | 4.84±0.89ac  |
| Heterotrophy     | Gln        | 16.97±1.41Aa  | 17.23±1.90Bb | 18.84±0.94Bb | 20.36±1.81Bb | 10.36±0.40Aa | 5.22±0.26Ad  | 20.36±1.81Bb |
|                  | Asn        | 1.16±0.22bc   | 2.80±0.18gb  | 3.31±0.50gb  | 2.61±0.44gb  | 0.95±0.21bc  | 0.67±0.29kc  | 2.61±0.44gb  |
|                  | Gln        | 0.67±0.15Ab   | 1.80±0.38Ab  | 0.70±0.20Ab  | 1.62±0.55Ab  | 4.45±0.48bcb | 6.09±1.30Ab  | 1.62±0.55Ab  |

The superscript letters indicate the statistical significance, identical letters identify means that are not significantly different, different letters identify statistically different means (P < 0.05). Upper case letters indicate the comparison between autotrophic and heterotrophic conditions.
Conclusions

In conclusion, the pigment and amino acid contents in *Galdieria sulphuraria* were profoundly dependent on the nitrogen status of the cells. Relevant differences were found between cells cultured in autotrophy or in heterotrophy, where the energy required for growth and metabolism depended on photosynthesis, or only on the respiration of added organic substrate, as in heterotrophic cells.

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