Biofilm Formation and Indole-3-Acetic Acid Production by Two Rhizospheric Unicellular Cyanobacteria

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Introduction

Agricultural soil provides an excellent ecological niche for microbial diversity governed by the soil and type of vegetation [14]. In particular, the rhizosphere, soil close to plant roots, is considered to be an environment rich in substrates for microbes, particularly for those that are associated with the roots of plants, such as various heterotrophic bacteria and fungi, as well as phototrophs like cyanobacteria and microalgae [9]. Rhizospheric microbes actively respond to the variety of metabolites released by plant roots. These microbes along with their metabolites also interact with plant roots in a variety of beneficial, harmful, and harmless ways [32, 33]. These interactions can control not only plant growth and development but also alter the plant's susceptibility to disease and abiotic stress [33]. In return, the plants modify the rhizospheric environment by providing the microorganisms with substrate through root exudates [28]. Plants secrete more than 40% of their photosynthates in the form of root

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Microorganisms that live in the rhizosphere play a pivotal role in the functioning and maintenance of soil ecosystems. The study of rhizospheric cyanobacteria has been hampered by the difficulty to culture and maintain them in the laboratory. The present work investigated the production of the plant hormone indole-3-acetic acid (IAA) and the potential of biofilm formation on the rhizoplane of pea plants by two cyanobacterial strains, isolated from rice rhizosphere. The unicellular cyanobacteria Chroococcidiopsis sp. MMG-5 and Synechocystis sp. MMG-8 that were isolated from a rice rhizosphere, were investigated. Production of IAA by Chroococcidiopsis sp. MMG-5 and Synechocystis sp. MMG-8 was measured under experimental conditions (pH and light). The bioactivity of the cyanobacterial auxin was demonstrated through the alteration of the rooting pattern of Pisum sativum seedlings. The increase in the concentration of L-tryptophan and the time that this amino acid was present in the medium resulted in a significant enhancement of the synthesis of IAA (r>0.900 at p=0.01). There was also a significant correlation between the concentration of IAA in the supernatant of the cyanobacteria cultures and the root length and number of the pea seedlings. Observations made by confocal laser scanning microscopy revealed the presence of cyanobacteria on the surface of the roots and also provided evidence for the penetration of the cyanobacteria in the endorhizosphere. We show that the synthesis of IAA by Chroococcidiopsis sp. MMG-5 and Synechocystis sp. MMG-8 occurs under different environmental conditions and that the auxin is important for the development of the seedling roots and for establishing an intimate symbiosis between cyanobacteria and host plants.

Keywords: Auxin, bioassay, Chroococcidiopsis sp., cyanobacterial biofilms, indole-3-acetic acid, Pisum sativum, plant hormone, Synechocystis sp.
exudates that can be actively released from the root (including mucilage) and passively released (diffusive compounds) owing to osmotic differences between the soil solution and the cell [33, 42]. Soil microbes are sensitive to changes in the environmental conditions that prevail in the rhizosphere. Cyanobacteria are well-known for their capacity to produce a variety of secondary metabolites of which many are infochemicals [22]. Many cyanobacterial secondary metabolites influence plant growth and development [24]. Cyanobacteria have been reported to give a benefit to plants by producing growth-promoting substances (resembling auxin and gibberellin), vitamins, polysaccharides, and antibacterial and antifungal compounds. Production of phytohormones is not limited to plants. Certain microbes, including cyanobacteria, produce all known phytohormones [42, 44]. The capacity of microbes to synthesize secondary metabolites such as auxins varies with minor fluctuations of environmental conditions in the surroundings [10]. The microbes adapt to the rhizospheric conditions to sustain their growth and maintain their metabolic activity [7]. Auxin-producing cyanobacteria have been reported to stimulate plant growth in vitro [24], in a hydroponics system [31], as well as under field conditions [23]. Surprisingly, although cyanobacteria are phototrophic organisms, they are able to inhabit and live in the dark rhizosphere. Root colonization of cyanobacteria is not limited to the ectorhizosphere (the soil immediately adjacent to the root) and rhizoplane (root epidermal and mucilage layer) but also extends to the endorhizosphere (root tissue including the endodermis and cortical layers) [2]. Some cyanobacteria penetrate the root cells and grow intracellularly [38]. The effectiveness of rhizospheric microbe at promoting growth depends on its density in the rhizosphere, and once a critical microbial density is reached then the biofilm as a whole acts as a plant growth-promoting unit [33]. Furthermore, owing to their extracellular polymeric coating, cyanobacteria have the ability to attach to solid substrates or to other organisms and form biofilms [26]. *Pisum sativum* is a dicot, commonly used as an experimental plant in many microbiological, physiological, genetical, and ecological studies [41, 43]. Its seedlings with hairy roots are used in bioassays with phytohormones [5, 19]. Thus, cyanobacteria that have the ability to produce auxin and form biofilms on roots may be interesting for biotechnological application in agriculture. The purpose of this study was to assess the capability of two unicellular rhizospheric cyanobacteria to form biofilms on seedling roots and to synthesize the plant hormone auxin under relevant environmental conditions (such as low pH and less light).

### Materials and Methods

#### Strains and Growth Conditions

The unicellular cyanobacteria *Chroococcidiopsis* sp. MMG-5 and *Synechocystis* sp. MMG-8 have been isolated from rice rhizosphere in the agricultural fields of the University of Punjab, Lahore, Pakistan (31°30′13″ N, 74°17′47.05″ E), in September 2006, using standard cyanobacterial isolation and purification techniques [2]. The isolates were characterized by 16S rRNA gene sequencing technique, and the GenBank accession numbers are FJ839355 and FJ839359, respectively. Axenic cultures were maintained in BG-11 [39] medium, kept under constant conditions with light (~150 µE/m²/s) provided by four 18W fluorescent tubes at a 16:8 light:dark cycle and the temperature maintained at 26 ± 2°C.

#### Morphology

The cyanobacteria were observed by light microscopy and confocal laser scanning microscopy (CLSM) following the methods described previously [2]. The cyanobacteria were dispersed in warm liquid agarose (1% (w/v)) on an object glass and covered with a cover slip and observed with a light microscope (Zeiss Axiosph, Oberkochen, Germany) using differential interference contrast (DIC) microscopy. CLSM was performed using a TCS-NT microscope (Leica, Heidelberg, Germany) equipped with an Argon-Krypton laser. Autofluorescence from cyanobacterial chlorophyll *a* and phycobiliproteins was used for observation without staining. The organisms were excited by wavelength 488 nm, and emitted wavelengths were collected with a 590 nm long-pass filter.

#### Determination of Chlorophyll *a* Concentration

For the estimation of growth, chlorophyll *a* was determined following the method of Tandeau de Marsac and Houmard [46]. Harvesting of the cyanobacteria was done by centrifugation at 10,000 × g for 10 min at 4°C. The pellets were extracted by 80% methanol for 2 h in the dark at 4°C. The extract was centrifuged at 10,000 × g for 10 min at 4°C, and the absorbance of the supernatant was measured spectrophotometrically at 665 nm against 80% methanol. The chlorophyll content (µg/ml) was calculated from the absorption at 665 nm (OD₆₆₅ nm) × 13.9.

#### Extraction of IAA

Released and intracellular auxins were extracted from the spent culture media and from the cyanobacterial cells, respectively, as reported previously [3, 42]. Briefly, 3-week-old cultures (incubated with 1,000 µg/ml of L-tryptophan supplementation) were harvested by centrifugation at 10,000 × g for 20 min at 4°C. For estimation of extracellular IAA, the pH of the supernatant was adjusted to 2.8 (using 1.0 M HCl) and extracted three times with three volumes of ethyl acetate. The extracts were evaporated at 37°C using a rotary evaporator (Heidelberg LABOROTA, Cole-Parmer, IL, USA) and the aqueous fraction during extraction was adjusted to pH 7.0 (using 1.0 N NaOH) and extracted three times with water-saturated n-butanol, followed by drying in a rotary evaporator.
The extracts obtained were filtered through membrane filters (Millipore, 0.45 μm). For intracellular IAA, the cyanobacterial cells were homogenized in liquid nitrogen using a mortar and pestle. IAA was extracted overnight at 4°C in 80% methanol containing 10 mg/l butylated hydroxytoluene as the antioxidant. The methanolic fraction was centrifuged at 5,000 × g for 10 min at 4°C, followed by filtering through filters (nominal pore size 1.2 μm; GF/C, Whatman). The filtrate was partitioned with ethyl acetate and water-saturated n-butanol as described above. The organic phases were dried under a stream of nitrogen gas and the residues were re-dissolved in methanol. All extracts were stored at -18°C and assayed within 48 h.

**Determination of Concentration of IAA**

IAA was measured by the Salkowski colorimetric method [18]. Cyanobacteria were grown in BG-11 medium supplemented with L-tryptophan (500 μg/ml). After 18 days of incubation, the cultures were harvested by centrifugation at 10,000 × g for 10 min at 4°C. The spent medium was filter-sterilized (Millipore filter, 0.45 μm) and mixed with two parts of Salkowski reagent (1 ml of 0.05 M FeCl₃ mixed with 50 ml of 35% HClO₃), and then incubated for 30 min in the dark at room temperature. The appearance of red color indicated the presence of an IAA. The absorption was measured spectrophotometrically at 535 nm against a control of 1 ml culture medium and 2 ml of Salkowski reagent. A standard curve was made from a concentration series of indole-3-acetic acid (3-indoleacetic acid; Sigma-Aldrich) solutions. All measurements were done in triplicate.

**Effect of L-Tryptophan on IAA Production**

IAA was determined in the culture medium at regular intervals during 5 weeks as described previously [3]. Each strain of cyanobacterium was inoculated in six flasks containing 100 ml of BG-11 medium supplemented with L-tryptophan (250, 500, 750, 1,000, 1,250, and 1,500 μg/ml). Another flask was used as the control and was not inoculated. Every week, cells from one flask were harvested for the estimation of cyanobacterial growth (chlorophyll a) and for the measurement of released IAA. Each experiment was performed in triplicate.

**Effect of pH on IAA Exudation**

The effect of pH on the synthesis and exudation of IAA by *Chroococcidiopsis* sp. MMG-5 and *Synechocystis* sp. MMG-8 was measured. For each of the two strains, five flasks were inoculated, containing 100 ml of BG-11 medium supplemented with 500 μg/ml L-tryptophan but adjusted to different pH values (5, 6, 7, 8, and 9) by adding HCl or NaOH. After three weeks of growth, the cultures were harvested and IAA was measured [3].

**Effect of Light-Dark Regime on IAA Production**

*Chroococcidiopsis* sp. MMG-5 and *Synechocystis* sp. MMG-8 were inoculated into 100 ml of BG-11 medium supplemented with 500 μg/ml L-tryptophan. Cultures were grown for three weeks under four light-dark regimes: 16:8, 8:16, 24:0, and 0:24 h. Cultures were harvested and IAA was measured from the spent media [3].

**Effect of Nitrate on IAA Production**

The strains MMG-5 and MMG-8 were grown in BG-11 medium supplemented with 500 μg/ml L-tryptophan in two sets. The first set contained 1.5 g/l NaNO₃ whereas the second set lacked NaNO₃. Cultures were harvested after three weeks of growth and IAA was measured in the spent media.

**Quantification of IAA by GC-MS**

*Chroococcidiopsis* sp. MMG-5 and *Synechocystis* sp. MMG-8 were grown for three weeks in BG-11 medium supplemented with 500 μg/ml L-tryptophan, after which 5 μg/ml of 5-methoxy-indole-3-acetic acid (5-Me-IAA; Sigma-Aldrich) was added to the cultures as an internal standard. Both intracellular and extracellular IAs were extracted and analyzed by GC-MS following the method of Gutierrez et al. [20].

**Root Bioassay**

The IAA bioactivity of both cyanobacterial strains was demonstrated by assessing the effect of culture supernatant on the root growth of pea (*P. sativum* var. Climax), as described previously [1, 3]. Pea seeds were obtained from the Punjab Seed Corporation, Lahore, Pakistan. Seeds were surface sterilized by washing them for 5 min in 0.1% HgCl₂ followed by repeated washing with sterile distilled water. Subsequently, the seeds (7–8 seeds) were placed in Petri dishes containing two layers of Whatman filter paper, which served to retain the moisture. Cultures of strains MMG-5 and MMG-8 incubated with 1,000 μg/ml of L-tryptophan were centrifuged at 10,000 × g for 10 min in sterile tubes. Filter-sterilized (Millipore filter, 0.45 μm) supernatant (1, 2.5, and 5 ml) was added to Petri dishes along with MilliQ water (Millipore Corporation, MA, USA) to make a total volume of 10 ml. The Petri dishes were kept in the dark for germination of the pea seeds. After germination the dishes were transferred to a 16:8 light:dark (200 μE/m²/s) regime. The number and length of the roots of the seedlings were determined in triplicate after 10 days. These measurements were repeated with supplementation of filter-sterilized supernatant of the cyanobacteria from cultures of different age (1–6 weeks). IAA was measured colorimetrically with the Salkowski reagent from the spent media at each measurement.

**Biofilm Formation**

Pea seeds were germinated in Petri dishes under axenic conditions for 10 days (16:8 h light:dark period, and light intensity of 200 μE/m²/s). Subsequently, the seedlings were suspended in 50 ml tubes with their roots immersed in 25 ml of 10-fold diluted BG11 medium. Three-week-old cyanobacteria cultures were harvested by centrifugation at 10,000 × g for 10 min and suspended in 10 ml of sterile MilliQ water (Millipore Corporation, MA, USA). Cyanobacteria were added to the suspension of seedling roots to a
final amount of 2 µg chl-a/ml. Tubes were incubated at 25°C at a 16:8 light–dark cycle (200 µE/m²/s light intensity) for 7 days to allow for biofilm formation on the seedling roots. Subsequently, the seedling roots were excised, and loosely attached cyanobacterial cells were removed by washing with sterile MilliQ water by squeeze bottle. The presence of cyanobacterial biofilm was observed by CLSM [2].

Confocal Laser Scanning Microscopy

CLSM of the roots and the image analysis were done as previously described [2, 4]. The root cells were stained overnight with the fluorescent dye 5-(4,6-dichlorotiazinyl) aminofluoroes (DTAF; Cat. # D-16; Invitrogen Corporations, USA) (1 mM). Excess stain was removed by washing twice with phosphate buffer saline, pH 8 (1×), followed by two washings with 0.1 M carbonate buffer, pH 9. CLSM was performed using a TCS-NT microscope (Leica, Heidelberg, Germany) equipped with an Argon-Krypton laser. For simultaneous imaging of emission fluorescence from DTAF and autofluorescence of chlorophyll a and phycobiliproteins, root sections were excited by 488 nm. Emitted wavelengths from DTAF were collected using band-pass filter 530/30 and autofluorescence with a 590-nm-long pass filter.

Results

Strain Characteristics

The two strains of cyanobacteria grew differently in BG-11 medium. Chroococcidiopsis sp. MMG-5 formed aggregates (Fig. 1A), while Synechocystis sp. MMG-8 grew homogenously (Fig. 1B).

Effects of Culture Age and L-Tryptophan Concentration on IAA Production

Colorimetric analysis of the cyanobacterial culture supernatants revealed that both tested parameters (i.e., culture age and initial L-tryptophan concentration) in media significantly (ANOVA, p < 0.05) affected IAA production by the two cyanobacteria. The estimated IAA increased with culture age (r = 0.986, p = 0.01) and the concentration of the precursor L-tryptophan (r = 0.955 – 0.977, p = 0.01) (Tables 1 and 2). In both cyanobacteria, the highest concentration of IAA was detected after 5 weeks of growth and with 1.5 mg/ml of L-tryptophan, amounting to 38.2 and 66.0 µg/g chl-a of IAA in Chroococcidiopsis sp. MMG-5 and Synechocystis sp. MMG-8, respectively. The IAA production rate depended on the initial L-tryptophan concentration in the media. In media with ≤500 µg/g chl-a of L-tryptophan supplementation, the IAA production rate was up to 6 µg/g chl-a/week, whereas in media with >500 µg/g chl-a of L-tryptophan, the IAA production rate was higher than 14 µg/g chl-a/week after 1 week of incubation.

Intra- and Extracellular IAAs

Both strains of cyanobacteria accumulated IAA intracellularly and released it in the medium during the 5 weeks of the experiment and showed a similar pattern of gradual increase in released and intracellular IAA with culture age (Figs. 2A and 2B). The maximum extracellular IAA was 60 µg/g chl-a in Synechocystis sp. MMG-8 after 5 weeks of cultivation, which was 33% higher than the extracellular IAA produced by strain MMG-5 (45 µg/g chl-a) during the same cultivation time. The intracellular IAA concentration was in the same order of magnitude in both cyanobacteria. The maximum intracellular IAA in MMG-8 cultures of 5 weeks was up to 31 µg/g chl-a, 14% higher than the 27 µg/g chl-a that was in strain MMG-5 of the same age.
Effect of pH on IAA Synthesis

The pH of the medium significantly affected the synthesis of IAA. IAA production in both cyanobacteria was enhanced by an acidic pH. *Chroococcidiopsis* sp. MMG-5 produced the highest amount of IAA (up to 43 µg/g chl-a) at pH 6, whereas *Synechocystis* sp. MMG-8 exuded the highest amount of IAA (more than 45 µg/g chl-a) at pH 5. Quantities of IAA decreased gradually with an increase in media pH up to 9 (Table 3).

Effect of Light Regime on IAA Synthesis

The light:dark period significantly (ANOVA, *p* < 0.05) affected IAA production by the two cyanobacteria. The amount of IAA produced in cultures grown under 8:16h light:dark was significantly higher than in cultures grown under any of the other three light regimes (continuous light, continuous dark, and 16:8h light:dark). The maximum amount of IAA was up to 40 µg/g chl-a for both

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**Table 1.** Time course of auxin concentration in *Chroococcidiopsis* sp. MMG-5 cultures with different amounts of tryptophan.

<table>
<thead>
<tr>
<th>Time (weeks)</th>
<th>Initial tryptophan concentration (µg/g chl-a)</th>
<th>Correlation coefficients (r)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>250</td>
<td>500</td>
</tr>
<tr>
<td>1</td>
<td>1.58 ± 0.055</td>
<td>3.87 ± 0.058</td>
</tr>
<tr>
<td>2</td>
<td>3.67 ± 0.053</td>
<td>8.32 ± 0.165</td>
</tr>
<tr>
<td>3</td>
<td>4.86 ± 0.073</td>
<td>14.3 ± 0.228</td>
</tr>
<tr>
<td>4</td>
<td>5.81 ± 0.065</td>
<td>16.12 ± 0.264</td>
</tr>
<tr>
<td>5</td>
<td>7.37 ± 0.060</td>
<td>19.75 ± 0.170</td>
</tr>
</tbody>
</table>

Correlation coefficients (r) of incubation time with given initial tryptophan concentration.

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**Table 2.** Time course of auxin concentration in *Synechocystis* sp. MMG-8 cultures with different amounts of tryptophan.

<table>
<thead>
<tr>
<th>Time (Weeks)</th>
<th>Initial tryptophan concentration (µg/g chl-a)</th>
<th>Correlation coefficients (r)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>250</td>
<td>500</td>
</tr>
<tr>
<td>1</td>
<td>3.87 ± 0.136</td>
<td>5.48 ± 0.121</td>
</tr>
<tr>
<td>2</td>
<td>7.66 ± 0.066</td>
<td>9.77 ± 0.078</td>
</tr>
<tr>
<td>3</td>
<td>9.77 ± 0.057</td>
<td>16.54 ± 0.033</td>
</tr>
<tr>
<td>4</td>
<td>13.27 ± 0.052</td>
<td>23.52 ± 0.034</td>
</tr>
<tr>
<td>5</td>
<td>14.52 ± 0.063</td>
<td>28.38 ± 0.033</td>
</tr>
</tbody>
</table>

Correlation coefficients (r) of initial tryptophan concentration with given incubation time.

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**Fig. 2.** Intra- and extracellular IAAAs.

Mean values of three replicates. Different letters indicate significant difference between treatments, using Duncan’s multiple range test (*p* = 0.05).
Chroococcidiopsis sp. MMG-5 and Synechocystis sp. MMG-8 grown under 8:16 h light:dark. There was a marginal difference between the production of IAA by cultures grown under 16:8 h light:dark and cultures grown under continuous light. The cyanobacteria did not grow in the dark and, consequently, no IAA was produced (Table 3).

**Effect of Nitrate on IAA Synthesis**

*Chroococcidiopsis* sp. MMG-5 could grow without combined nitrogen and produced IAA under these conditions. IAA production was significantly higher (ANOVA, p < 0.05) (28%) in medium containing nitrate. *Synechocystis* sp. MMG-8 was unable to grow without combined nitrogen (Table 3).

**Quantification of IAA by GC-MS**

Extracts of intracellular and extracellular auxins were analyzed by using GC-MS in order to verify and quantify the IAA. The results confirmed the presence of intra- and extracellular IAAs in both strains. The amount of extracellular IAA in strain MMG-8 was 17% higher than in strain MMG-5 (Fig. 3). There was no difference in intracellular IAA between both strains.

**Effect of Supernatant on Root Length and Number of Lateral Roots**

Cyanobacterial supernatant containing high concentrations of IAA gave a negative effect on root length when compared with sterile growth medium. Five milliliters of medium supernatant of *Chroococcidiopsis* sp. MMG-5 containing 1.58 µg/g chl-a of IAA resulted in an increase of root length (up to 60%) in one week, as compared with the non-amended control. The increase of root length became less with age of the culture and disappeared after 5 weeks, while after 6 weeks of growth root length started to decrease (r = -0.995, p = 0.01). Treatment with 1 ml of supernatant resulted in a gradual increase in root length with culture age (i.e., increase in IAA) with a significant positive correlation of 0.974 at p = 0.01 (Fig. 4A). When using 1 ml of supernatant of *Synechocystis* sp. MMG-8, root length increased with a maximum of 81% after 3 weeks of growth, whereas 2.5 and 5 ml did not result in an increase of root length, and after 5 weeks of growth the roots became even shorter than in the control (Fig. 4B).

The number of lateral roots increased when cyanobacterial culture supernatant was added, and a significant positive correlation between the number of lateral roots and the concentration of IAA was found. Five milliliters of *Chroococcidiopsis* sp. MMG-5 cultural supernatant after 5 and 6 weeks of growth resulted in the highest increase in the number of roots (156% and 140%, respectively), as compared with the control (Fig. 5A). *Synechocystis* sp. MMG-8 also increased the number of roots with culture age and the volume of supernatant added. The highest stimulation was with 5 ml of supernatant, increasing 149%, 152%, 192%, and 167% after 3, 4, 5, and 6 weeks of growth, respectively. The correlation of IAA concentration and root number was highly significant (r = 0.995, p = 0.01) (Fig. 5B).

**Table 3. Effect of growth conditions (pH, light:dark period, and nitrates) on auxin biosynthesis (µg/g chl-a) by cyanobacterial strains.**

<table>
<thead>
<tr>
<th>Strains</th>
<th>pH</th>
<th>Light</th>
<th>Nitrates</th>
<th>16:08</th>
<th>8:16</th>
<th>CL</th>
<th>CD</th>
<th>N+</th>
<th>N-</th>
</tr>
</thead>
<tbody>
<tr>
<td>MMG-5</td>
<td>39</td>
<td>43</td>
<td>30</td>
<td>14</td>
<td>6</td>
<td>33</td>
<td>40</td>
<td>34</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>± 2.12(e)</td>
<td>± 2.36(f)</td>
<td>± 1.24(d)</td>
<td>± 1.17(b)</td>
<td>± 0.77(a)</td>
<td>± 0.83(b)</td>
<td>± 1.54(d)</td>
<td>± 2.33(b)</td>
<td>± 0.00(a)</td>
</tr>
<tr>
<td>MMG-8</td>
<td>45</td>
<td>37</td>
<td>30</td>
<td>20</td>
<td>5</td>
<td>37</td>
<td>42</td>
<td>35</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>± 1.73(f)</td>
<td>± 2.43(e)</td>
<td>± 1.92(d)</td>
<td>± 1.02(c)</td>
<td>± 0.85(a)</td>
<td>± 1.22(cd)</td>
<td>± 1.77(d)</td>
<td>± 1.26(b)</td>
<td>± 0.00(a)</td>
</tr>
</tbody>
</table>

16:08, 8:16 = light:dark period in hours; CL = complete light; CD = complete dark.
Mean values of three replicates. Different letters indicate significant difference between growth conditions, using Duncan’s multiple range test (p < 0.05).

Fig. 3. Comparison of intra- and extracellular IAAs in *Chroococcidiopsis* sp. MMG-5 and *Synechocystis* sp. MMG-8 as analyzed by GC-MS.
Formation of Biofilms on the Rhizoplane of *Pisum sativum* Seedlings

The cyanobacteria showed ability to colonize the rhizoplane. *Chroococcidiopsis* sp. MMG-5 extensively colonized the root surface and concentrated in the grooves between adjacent surface cells (Figs. 6A and 6B). *Synechocystis* sp. MMG-8 was scattered on the root surface but sometimes formed colonies (Figs. 6C and 6D). Furthermore, the orthogonal sectioning by CLSM illustrates how the cyanobacteria invaded the epidermis and the cortex region, because they appeared inside the root cells or underneath the outermost layer. *Chroococcidiopsis* sp. MMG-5 was mainly present at a depth of 10–15 µm inside the root cells (Fig. 7A). *Synechocystis* penetrated the plant cells to a depth of 5–7 µm (Fig. 7B).

Discussion

This study shows the ability of rhizospheric unicellular cyanobacteria to produce IAA and to colonize the roots, forming a thick biofilm. These two traits are considered to be pivotal for rhizospheric microbes, and any microorganism possessing these abilities should be successful in agricultural ecosystems [35].

Cyanobacteria have the ability to attach to surfaces through their extracellular polysaccharides. Such cyanobacteria are considered as important colonizers of surfaces and initiate the development of biofilms [17]. In order to allow the establishment of an intimate association of rhizospheric microorganisms, the plant root surface acts as the substrate to which microbes attach. The roots assist in this process by
exuding a variety of signal compounds that attract the soil microorganisms [6]. The formation of a biofilm on the roots is an important trait of rhizospheric microorganisms, which prevents them from being dissociated from the plant caused by various chemical, biological, or physical processes in the soil [13]. Colonization of microbes on the plant root surface is not uniform, but instead occurs in patches along the root, which ultimately covers ~15–40% of the total plant root surface [32]. The formation of a biofilm depends on several factors such as the number of attached cells per area of root surface, ability to withstand abrasive forces, competition with other microorganisms, and the potential to penetrate the root cells. Both of the investigated cyanobacteria colonized the rhizoplane. Once a biofilm formed on the rhizoplane, the cyanobacteria started to penetrate the root and colonized intracellular spaces or even penetrated the root cells or endorhizosphere (root tissue including the endodermis and cortical layers), probably in order to establish a more intimate association. It is not clear what the advantage of the penetration of the root cells is, nor what the mechanism behind it is. It has been suggested that proteolytic enzymes play a role in the degradation of the plant cell wall, thereby allowing the organism to enter the cell, but it could also be achieved by mechanical force [2, 34]. Furthermore, the extracellular auxins have been reported to up-regulate the production of cellulases and hemicellulases in plants, which ultimately leads to the disintegration of the cell wall. Such weak patches might be used by microbes to penetrate the plant roots [20].

The most common pathway for IAA biosynthesis in plants and microorganisms is from L-tryptophan [42]. Both strains of cyanobacteria produced IAA only in the presence of L-tryptophan. Under natural conditions, the plant roots are the source of L-tryptophan, provided in the form of root exudates that serve as the substrate for the rhizospheric microorganisms. Cyanobacteria have been reported to synthesize IAA from L-tryptophan, and it depends on its concentration as well as culture age [3, 42]. Synechocystis sp. MMG-8 converted up to 4.5% of L-tryptophan to IAA. The metabolism of cyanobacteria is sensitive to environmental factors such as pH and light [37]. This became apparent by the effect of pH and light on the IAA biosynthesis. Generally, cyanobacteria prefer neutral or alkaline pH for growth [38], but low pH appeared to be favorable for optimal auxin production. It is well-established that the rhizosphere is generally characterized by a low pH (pH 5.5), and many microbial genes involved in the interaction with plants are expressed higher at this pH. The key gene for auxin biosynthesis (ipdC) is highly expressed at low pH [16, 40, 48]. The light:dark regime is an important factor for cyanobacteria, and the metabolic activities are generally stimulated with the increase of the photoperiod [45]; A considerable difference was observed in auxin biosynthesis by the cyanobacteria incubated under various light:dark regimes. Maximum IAA was detected under a 8:16 h light:dark regime. Light is directly involved in controlling the auxin levels in plants by regulating the auxin synthesis genes and translocation of auxins throughout the plants [21]. Light regulates the genes that code for enzymes involved in auxin synthesis and proteins involved in auxin translocation such as PIN-formed proteins (PIN) and P-glycoprotein (PGP) [21]. Genes whose products are involved in tryptophan-dependent IAA biosynthesis are especially under control by environmental light. By positive and negative regulation of auxin biosynthesis genes, the plant maintains a certain auxin concentration, according to what its developmental stage requires [30, 47]. Auxins, and especially IAA, are sensitive to light and are readily degraded through photolysis [15, 27]. Compared with light:dark cycles with long dark periods, the relatively low amounts of IAA that were found after the long exposure of the cyanobacteria to light might have been the result of photolysis of IAA.

Fig. 6. Rhizoplane colonization by cyanobacteria (A and B) Chroococcidiopsis sp. MMG-5 and (C and D) Synechocystis sp. MMG-8. Images obtained by a confocal laser scanning microscope. Green shows plant root cells stained with DTAF. Red shows cyanobacterial cells.
The possible reasons for the lower IAA yield in the absence of nitrate might be that (i) it restricted growth, (ii) tryptophan appeared to inhibit nitrogen fixation [36], and (iii) the fixation of nitrogen may have prevented the cells of synthesizing IAA, thus resulting in a low IAA in the supernatant. IAA measured by GC-MS gave significantly lower amounts than the colorimetric method (Salkowski’s reagent). The difference between the GC-MS technique and the colorimetric assay is attributed to the fact that the former only measures the amount of IAA, whereas the Salkowski reagent also reacts with other indolic compounds and therefore is a measure of all auxins present [20, 42].

Fig. 7. Display windows of a Z stack in XY horizontal CLSM sections of (A) *Chroococcidiopsis* sp. MMG-5 and (B) *Synechocystis* sp. MMG-8 (red) with seedling roots (green). Each slice represents an image at a different depth above the cell surface.
Hence, both methods do not measure the same, and the results indicate that other auxins than IAA are produced and may be involved in the observed effects on the root development.

Like other phytohormones, auxins are a group of important signaling molecules involved in the interaction between rhizospheric microbes and plants that affect plants even at very low concentrations (10^{-5}–10^{-6} M) [11, 25]. Roots are the most sensitive parts of plants with respect to the effect of auxins. Hence, root length and the number of lateral roots have been used as a bioassay for auxins, particularly for IAA [8]. Different volumes of sterile supernatant from the growth medium of cyanobacterial cultures of different age were used in the root bioassay in order to detect IAA-like activity. Pea (P. sativum) was used as the test plant. The concentration of secreted IAA in the culture supernatant increased with the culture age. A gradual change in the root parameters (root length and number of lateral roots) was observed with the concentration of IAA. The concentration of auxin in or near roots is critical because a slight change results in a change in root length. On the one hand, IAA in low concentration causes an increase in root length; on the other hand, above a critical concentration, auxin inhibits the growth of the root or even causes a decrease of its length. This threshold concentration varies between different plants [12, 29]. Highly significant correlations were found between the concentration of IAA in the supernatant and the root parameters (root length and number of lateral roots). This could indicate an involvement of cyanobacterial IAA in the regulation of root growth [3]. Thus, the rhizospheric cyanobacteria showed useful traits (auxin synthesis and biofilm formation that possibly lead to intense root colonization) in vitro, as they do in nature. This ability of rhizospheric cyanobacteria can be exploited for agriculture.

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