DevT (Alr4674), resembling a Ser/Thr protein phosphatase, is essential for heterocyst function in the cyanobacterium *Anabaena* sp. PCC 7120

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Heterocyst-forming cyanobacteria are able to perform oxygenic photosynthesis and nitrogen fixation simultaneously in the same filament, by restricting the highly O2-sensitive nitrogenase to specialized cells, the heterocysts. A remarkable change in morphology and metabolism accompanies the differentiation of heterocysts, which only occurs when no source of combined nitrogen is available. In this study, we characterized DevT (Alr4674), a putative protein phosphatase from *Anabaena* PCC 7120. Mutants defective in devT are able to form morphologically mature heterocysts, which however cannot fix N2, and the mutant cannot survive without a source of combined nitrogen. DevT shows homology to phosphatases of the PPP family and displays a Mn2+-dependent phosphatase activity that can be inhibited by phosphatase inhibitors and oxidizing conditions. DevT is constitutively expressed in both vegetative cells and heterocysts, and is not regulated by NtcA. The heterocyst regulator HetR may exert a certain inhibition on the expression of devT. Under diazotrophic growth conditions, DevT protein accumulates specifically in mature heterocysts. Therefore DevT plays a still unknown role in a late essential step of heterocyst differentiation.

INTRODUCTION

Protein phosphorylation plays an important role in countless cellular processes, including regulation of enzyme activity, assembly of macromolecules and proteolysis. Most abundant in bacteria is the phosphorylation of proteins at specific His and Asp residues involved in so-called two-component systems. Those systems are based on phospho-relay signal-transduction pathways that allow the cells to respond to environmental changes. In the last two decades other types of phosphorylation that were thought to be restricted to eukaryotes, the phosphorylation of Ser/Thr and Tyr residues, have been recognized in bacteria (Shi et al., 1998). Many examples of these eukaryotic systems have been found in the increasing number of available bacterial genome sequences. In particular, many protein sequences homologous to Ser/Thr and Tyr kinases and phosphatases have been identified in the diverse group of cyanobacteria (Zhang et al., 1998b, 2005). However, the function of most of these enzymes is still largely unknown. The enormous quantity of two-component systems as well as protein kinases and phosphatases in strains able to undergo cellular differentiation in adaptation to environmental stress has led to the suggestion that protein phosphorylation is related to the cellular complexity of these cyanobacteria (Kennelly & Potts, 1996; Wang et al., 2002). Interestingly, the Ser/Thr kinases from the heterocyst-forming cyanobacterium *Anabaena* sp. PCC 7120 (hereafter referred to as *Anabaena*) exhibit very complex domain structures. In several cases regulatory domains like the GAF domain or His kinase domains are fused to the Ser/Thr kinase part of these enzymes. In addition, several genes

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Abbreviations: DAB, diaminobenzidine; pNPP, p-nitrophenyl phosphate; TTC, 2,3,5-triphenyltetrazolium chloride.
The GenBank/EMBL/DDBJ accession number for the devT (alr4674) gene is CAB57788.
Two supplementary figures are available with the online version of this paper.
encoding Ser/Thr kinases or phosphatases are clustered with genes for proteins involved in two-component systems, and these protein-modification systems may be part of the same signalling pathway (Wang et al., 2002; Zhang, 1996; Zhang et al., 1998b).

Ser/Thr phosphatases are the counterparts of Ser/Thr kinases and can be divided into two main families, the PPP and the PPM family. Phosphatases of the PPP family are the most frequent eukaryotic Ser/Thr protein phosphatases. Examples are protein phosphatases PP1, PP-2A and PP-2B (calcineurin), which exhibit invariant residues involved in catalysis (Barton et al., 1994). They utilize the same binuclear metal-ion assisted catalytic mechanism, but have strikingly different structures (Barford, 1996). Crystal structures have been determined for PP-1 and PP-2B, showing the conserved metal-binding sites in the catalytic centre: Mn$^{2+}$ and Fe$^{2+}$ in PP-1, Zn$^{2+}$ and Fe$^{2+}$ in PP-2B (reviewed by Cohen, 1989; Shi, 2004). Less-conserved regulatory subunits determine activity, substrate specificity and function (Cohen, 1989). It is not known whether prokaryotic phosphatases of the PPP family show similar distribution into non-catalytic and catalytic domains. The second family of Ser/Thr protein phosphatases is the PPM family with the representative phosphatase PP2C, which differs clearly from the PPP-type in primary sequence, structure and catalytic mechanism (Mumby & Walter, 1993; Virshup & Shenolikar, 2009).

Several filamentous cyanobacteria, such as members of the genera Nostoc and Anabaena, are able to simultaneously assimilate CO$_2$ and N$_2$ and to perform oxygenic photosynthesis in one multicellular filament, despite the fact that the nitrogenase, the enzyme responsible for nitrogen fixation, is very sensitive to oxygen. A sophisticated solution to this problem is realized by some filamentous diazotrophic cyanobacteria: they restrict the oxygen-sensitive nitrogen-fixation process to specialized cells termed heterocysts, neighbouring vegetative cells. The fixation process is adapted to the vegetative cells, and nitrogenase is repressed in the differentiating cells, whereas expression of genes required for morphological changes or nitrogen fixation is induced (Wolk et al., 1994). The global nitrogen regulator NtcA plays a key role in these regulatory processes as either activator or repressor of several genes (Herrero et al., 2001, 2004; Zhang et al., 2006). The second key regulator is HetR, a protein exhibiting protease and DNA-binding activities in vitro, which regulates expression of genes required for heterocyst differentiation, in some cases indirectly via activation of NtcA expression (Black et al., 1993; Buikema & Haselkorn, 1991; Flores & Herrero, 2010; Huang et al., 2004; Khudyakov & Golden, 2004; Muro-Pastor et al., 2002; Zhou et al., 1998).

In addition to these key players, several protein kinases and phosphatases required for growth on N$_2$ as nitrogen source have been identified (Kumar et al., 2010; Ohmori et al., 2001; Zhang et al., 2005). The pknE gene of Anabaena encoding a Ser/Thr kinase clusters with the prpA gene encoding a protein phosphatase, and both are required for diazotrophic growth (Zhang et al., 1998a). The genes are upregulated after 8 h of nitrogen depletion (Ehira et al., 2003). Mutants in these genes show aberrant heterocyst envelope structures and are not able to restore nitrogenase activity after incubation under micro-oxic conditions (Zhang et al., 1998a). A mutant in the pknD gene, encoding another Ser/Thr protein kinase, differentiates normal heterocysts but shows reduced growth on N$_2$, even though the rate of nitrogen fixation is similar to that of the wild-type Anabaena (Zhang & Libs, 1998). This phenotype led to the suggestion that PknD might be involved in the trafficking of the fixed nitrogen from the heterocyst to the surrounding vegetative cells. The prpJ1 and prpJ2 genes, encoding PP2C-type protein phosphatases, were identified as new regulators involved in the initiation of heterocyst differentiation. The activity of PrpJ1 is essential during heterocyst maturation, regulating the synthesis of one of the two heterocyst glycolipid layers. The closely related phosphatase PrpJ2 was proposed to act together with PrpJ1 in upregulating hetR (Jang et al., 2007, 2009). In the present work, we describe a novel potential Ser/Thr protein phosphatase, encoded by the devT gene (alr4674), which is essential for growth on N$_2$ in the cyanobacterium Anabaena.

**METHODS**

**Bacterial strains and growth conditions.** Anabaena sp. strain PCC 7120 (Anabaena) and derived mutant strains were grown in liquid medium according to Allen & Arnon (1955) under photoautotrophic conditions as described previously (Fiedler et al., 1998b). Routinely, Anabaena strains were grown in Erlenmeyer flasks with constant shaking. Experimental liquid cultures were grown in 750 ml bottles, bubbled with 2% CO$_2$-enriched air. During nitrogen step-down experiments, liquid cultures were washed three times in medium without combined nitrogen and resuspended in the same medium to
induce heterocyst formation. Mutant s2 (Ernst et al., 1992), which is a transposon (Tn5-1065)-derived mutant in the devT gene (alr-6674) of Anabaena, strain DRpMA54 (a site-directed mutant of devT) and hetR mutant DR884a (Black et al., 1993) were grown in the presence of 5 mM NO₃⁻ and 50 µg neomycin ml⁻¹. Mutant s2 (pMA40) was grown in the presence of 50 µg neomycin and 5 µg spectinomycin ml⁻¹. ntcA mutant CSE2 (Frias et al., 1994), was grown in the presence of 5 mM NH₄Cl supplemented with streptomycin and spectinomycin, 2.5 µg ml⁻¹ each.

Strains of Escherichia coli were grown in LB medium containing appropriate antibiotics: kanamycin, ampicillin, spectinomycin (50 µg ml⁻¹), streptomycin or chloramphenicol (25 µg ml⁻¹) (Sambrook et al., 1989). Conjugation between E. coli and Anabaena was performed by tripontial mating using RP-4 as conjugal plasmid and pRL528 as helper plasmid (Elhai & Wolk, 1988b; Wolk et al., 1984).

**DNA isolation and manipulation.** To isolate plasmid DNA from E. coli, the Qiagen plasmid kits were used. DNA from Anabaena strains was isolated as described by Cai & Wolk (1990). Restriction digestions and ligations were carried out with enzymes and protocols from New England Biolabs, Fermentas, Eppendorf or Roche. Sequencing was performed with a T7 sequencing kit from Pharmacia using primers and ligations were carried out with enzymes and protocols from New England Biolabs, Fermentas, Eppendorf or Roche. 

Inactivation of the devT gene was done by cloning the C.K3 cassette, conferring neomycin/kanamycin resistance, flanked by Smal sites from the MCS of pRL448 (Elhai & Wolk, 1988a) into the unique Ndel site of pML71, which had been blunted by Klenow treatment, resulting in plasmid pMA53. A 4.1 kb insert was cloned from that plasmid as a HindIII fragment into Ndel-digested mobilizable suicide vector pRL277 (Black et al., 1993), resulting in plasmid pMA54. Transfer of pMA54 to wild-type cells of Anabaena by tripontial mating, followed by selection on neomycin plates and sucrose, resulted in clones in which the devT allele had been replaced by devT::C.K3 as verified by PCR (data not shown). One chosen clone was the double recombinant DRpMA54.

To carry out complementation of mutants s2 and DRpMA54, first the devT gene and its flanking regions were isolated as an 8 kb Clal fragment of plasmid pML71 and cloned into the Clal site of shuttle vector pRL1049 (Black & Wolk, 1994), resulting in plasmid pMA40. After transferring this plasmid to mutants s2 and DRpMA54 by tripontial mating, exconjugants were selected on spectinomycin and tested for their ability to grow on N₂ as nitrogen source.

To generate a DevT–GFP reporter protein, a DNA fragment consisting of 400 bp of the upstream region of devT plus its entire coding region was amplified by PCR using genomic DNA as template with primers DevTGFP1F (5'-AGTACGAATTCTCGGTTAAG-AATCTTTGGC-3') and DevTGFP1R (5'-CTATCCCGCCACAGGATGACTACTG-3'). The PCR product was cloned into the unique EcoRI site of plasmid pCSEL19 (Olmedo-Verd et al., 2006) in-frame with the ORF of gfp. The resulting plasmid, named pIM333, was used as a template for a second PCR with primers DevTGFP2F (5'-CGGGAGAATTCTCGGTTAAGAAATC-3') and DevTGFP2R (5'-TGCAGGCAGTCGATTGGATC-3'). The PCR product obtained was restricted with EcoRI/PstI and cloned into EcoRI/PstI-digested pCSEL24, designed for integration into the nucA region in the z-megaplasmid of Anabaena (Olmedo-Verd et al., 2006), rendering plasmid pIM333. Conjugation to wild-type Anabaena resulted in single recombinants, whose genomic structures were confirmed by PCR (data not shown).

**RNA isolation and devT expression analysis.** The protocol for RNA isolation for Northern blot analysis was described by Muro-Pastor et al. (1999). In Northern blots, 30 µg RNA per lane was used. The devT probe was a 371 bp PCR fragment generated with primers 230 (5'-GGTGGGATTTGTGAAATGTCG-3') and 231 (5'-GATTAATTGCTCAGAAGGTG-3') and labelled using a ReadyToGo DNA Labelling kit (Amersham) and [γ-32P]dCTP. The rnpB probe was used as a control for RNA loading, and was labelled as described by Muro-Pastor et al. (1999). After 2 h of prehybridization, the probe was hybridized overnight at 65 °C and the results were visualized with a Cyclone Storage Phosphor System and OptiQuant image analysis software (Packard).

To isolate RNA for RT-PCR analysis, 50 ml cells were harvested by centrifugation and RNA was isolated by hot-phenol pre-extraction and cell lysis combined with the High Pure RNA isolation kit (Roche) following the manufacturer’s recommendations.

The RT-PCR analysis of devT mRNA abundance under different growth conditions was performed by using the Qiagen OneStep RT-PCR kit according to the manufacturer’s description, with 20 ng total RNA per reaction as template. Primers used were oligo230 (5'-GGTGGGATTTGTGAAATGTCG-3') and oligo217 (5'-GGCCGATTCTGTGGCAG-3') for devT amplification, and oligo362 (5'-GGCTTTGCGGTGTCGACG-3') and oligo130 (5'-TAACCGGGGTTCTTTGCCTGTG-3') for rnpB. The mRNA level of the constitutively expressed rnpB gene was used as a loading control.

**Microscopy.** Electron microscopy was done as described by Fiedler et al. (1998). In brief, fixation and post-fixation was done with glutaraldehyde and potassium permanganate; ultrathin sections were stained with uranyl acetate and lead citrate. The samples were examined with a Zeiss EM109 electron microscope at 80 kV.

Bright-field micrographs were taken with a Leica microscope equipped with a DFC420 C camera. DAB (diaminobenzidine) staining was performed according to Ernst et al. (1992). To observe TTC (2,3,5-triphenyltetrazolium chloride) reduction, cells were supplied with TTC at final concentration of 0.01 % (w/v) for 30 min and subsequently fixed with 10 % glutaraldehyde and stored on ice in the dark until examined (Fay & Kulasooriya, 1972).

Accumulation of DevT–GFP fusion protein was analysed by fluorescence microscopy. Samples were observed using a Leica HCN PL APO 63 x/1.3 GLYC objective attached to a Leica DM5500B microscope connected to an external light source for fluorescence excitation Leica EL6000 (Leica, Wetzlar, Germany). The wavelength of excitation was 488 nm. Fluorescent emission was monitored by collection across windows of 500–570 nm (GFP imaging) and 630–700 nm (cyanobacterial autofluorescence).

**O₂ respiration and nitrogenase activity measurements.** To determine O₂ consumption, Anabaena cultures were grown to mid-exponential phase. The amount of cells equivalent to 20 µg chlorophyll a (measured in methanolic extracts as described by Mackinney, 1941) were collected and concentrated to 2 ml. Subsequently, cells were incubated in constant light (100 µmol photons m⁻² s⁻¹), and O₂ production was monitored using a Clark-type oxygen electrode (Hansatech Instruments), until it reached a plateau. The cultures were then immediately supplied with 40 µM DCMU, which inhibited O₂ production, and the O₂ consumption ratio was determined. Nitrogenase activity was measured as described by Valladares et al. (2007).

**Expression and purification of His-tagged DevT.** A construct of devT fused at its 3’ end to six codons encoding histidine was prepared as follows. PCR was performed with primers 38 (5'-GTGGGGATTTGTGAAATGTCG-3') and 39 (5'-TAACCGGGGTTCTTTGCCTGTG-3') and plasmid pML71 as template. The resulting
fragment contained the entire ORF of devT flanked by SpnI and BamHI sites. After digestion with both restriction enzymes, the devT fragment was ligated into the SpnI and BamHI sites of cloning vector pQE70 (Qiagen), resulting in plasmid pLM88. For overproduction of His$_6$-tagged DevT (in the following termed DevT-H$_6$), an overnight culture of E. coli M15 containing plasmids pREP4 (providing additional copies of the LacI repressor) and pLM88 was re inoculated 1:60 in 500 ml LB medium (supplemented with 100 μl ampicillin ml$^{-1}$ and 25 μl kanamycin ml$^{-1}$) and grown at 37 °C. After the culture had reached an OD$_{600}$ of 0.6, IPTG was added to a final concentration of 1 mM and incubation was continued for 4 h at 37 °C. After harvesting the cells by centrifugation, the cell pellet was resuspended in 1 ml per g cell pellet of lysis buffer (50 mM potassium phosphate buffer pH 7.8, 300 mM NaCl, 10 mM imidazole, 1 mg lysozyme ml$^{-1}$) and incubated on ice for 30 min. Cells were disrupted by sonication with a Sonifier B-12 from Branson Sonic Power Company. The soluble proteins of the crude extract were applied to a Ni$^{2+}$-NTA column (Qiagen), equilibrated with 50 mM phosphate buffer pH 7.8, 300 mM NaCl, 20 mM imidazole. The same buffer was used for washing. Elution was done with 50 mM phosphate buffer pH 7.8, 300 mM NaCl and 250 mM imidazole. After analysis by SDS-PAGE, the fractions containing pure DevT-H$_6$ were pooled and dialysed overnight against 10 mM potassium phosphate buffer pH 7.8, 1 mM EDTA, and stored at −20 °C until further use. All steps after crude extract preparation were performed at room temperature to prevent precipitation of DevT-H$_6$. Most of the expressed protein was found in inclusion bodies, although for purification only the soluble fraction was used. N-terminal sequencing of the purified protein was done by Rainer Deutzmann, Biochemie 1, Universität Regensburg, Germany.

Biochemical methods. The amount of protein was determined with the BCA kit from Pierce. SDS-PAGE was performed with the Laemmli buffer system using a Bio-Rad Minicell electrophoresis chamber. Ten micrograms of protein, denatured in SDS loading buffer for 5 min at 95 °C, was applied per lane. Broad-range SDS-PAGE standards from Bio-Rad were used as size markers. Coomassie blue was used to stain the proteins.

Size-exclusion chromatography was performed on a Superdex 200 10/300 GL column, equilibrated with gel filtration buffer (10 mM sodium phosphate buffer pH 7.0, 5 mM MgCl$_2$, 0.5 mM EDTA, 1 mM DTT and 400 mM NaCl). The column was loaded with 500 μg DevT-H$_6$, in 500 μl gel-filtration buffer and developed on a AKTA chromatography system (GE Healthcare) at a flow rate of 0.5 ml min$^{-1}$, with 0.5 ml fractions collected. Aliquots (20 μl) from each fraction were analysed by SDS-PAGE and Coomassie blue staining to show the presence of DevT-H$_6$. The Superdex column was calibrated by two independent 500 μl injections corresponding to a total of five proteins: cytochrome c (12.4 kDa), carbonic anhydrase (29 kDa), BSA (66 kDa), alcohol dehydrogenase (150 kDa) and β-amylase (200 kDa).

Standard phosphatase activity of DevT-H$_6$ was assayed in 1 ml volumes in a containing 10 mM Tris/HCl pH 8.2, 2 mM MnCl$_2$, 1 mM DTT, 50 mM NaCl and 5 mM p-nitrophenyl phosphate (pNPP) as substrate. The reaction was started by adding 1–10 μg purified DevT-H$_6$ and the increase in A$_{405}$ was measured for 15 min in a Kontron UVicron spectrophotometer. The same reaction omitting DevT-H$_6$ was prepared as a blank. To determine the optimum temperature for the DevT activity, the reaction was performed in test tubes in heating blocks, stopped after 15 min by chilling on ice and the A$_{405}$ was measured against the blank, incubated under the same conditions. To measure the pH dependence of DevT activity, the pH value was adjusted in the range from 7.4 to 8.6 with Tris/HCl present in the standard buffer. When used, phosphatase inhibitors were present at a final concentration of 10 mM. To measure the effect of H$_2$O$_2$ on DevT activity, DevT-H$_6$ was incubated in the presence of 0.3, 0.5, 1, 1.5 and 2 mM H$_2$O$_2$ in 10 mM HEPES, 1 mM EDTA for 30 min at 30 °C. The reaction was stopped by adding 100 units catalase and the specific activity determined after 10 min at 37 °C.

To determine the time-course of inhibition of DevT by H$_2$O$_2$, DevT-H$_6$ at a concentration of 1 μg ml$^{-1}$ was incubated with 1 mM H$_2$O$_2$ in 10 mM HEPES, 1 mM EDTA pH 8.0 at 30 °C. Every 5 min, 10 μl aliquots were removed and assayed by the pNPP standard test to measure the specific activity of DevT.

To test reactivation of DevT by DTT, 10 μg DevT-H$_6$ was first inactivated by 1 mM H$_2$O$_2$, in 10 mM HEPES, 1 mM EDTA pH 8.0 for 30 min at 30 °C. After removing H$_2$O$_2$ by addition of 100 units catalase, DevT activity was recovered by incubation with 10 mM DTT for 15 min at 37 °C. The activity was measured in the pNPP standard test against the control sample (DevT-H$_6$) treated in the same way.

Immunodetection of DevT protein. Antiserum against purified DevT-H$_6$ was produced in rabbits (Biogenes, Berlin). After 90 days of immunization, IgGs from the obtained antiserum were further purified by incubation with purified DevT-H$_6$ immobilized on a PVDF membrane.

To prepare soluble proteins from Anabaena, cells were resuspended in lysis buffer (20 mM HEPES pH 7.5, 1 mM PMSF, 1 mM DTT and 0.5 mM EDTA) and subjected to five 30 s rounds of sonication (45 % duty cycle, 50 % power) in a Branson sonifier 250. Samples were briefly centrifuged at 4 °C, 2 min, 1000 g and the supernatant was taken and used in Western blots. Isolation of heterocyst-specific proteins was carried out essentially as described by Moslavac et al. (2007). Isolated heterocysts were resuspended in lysis buffer and samples were passed through a French press (10 cycles at 1300 p.s.i., −9 MPa). Samples were centrifuged at 4 °C for 5 min at 15000 g and the supernatant was used in Western blot analysis.

For Western blots, proteins were resolved by SDS-PAGE and transferred to PVDF membranes using a semidyed transfer system (Peqlab Biotechnologies). Membranes were blocked in TBS (20 mM Tris pH 7.5, 150 mM NaCl) containing 5 % (w/v) BSA. Filters were incubated with purified anti-DevT antiserum in TBS containing 5 % (w/v) BSA, 5 % (v/v) IgG solution and incubated with horseradish peroxidase conjugate (1 : 15 000) (Amersham Biosciences). Immunoreactive bands were detected using the Lumi-Light Western blotting conjugate (1 : 15 000) (Amersham Biosciences). Membranes were blocked in TBS (20 mM Tris pH 7.5, 150 mM NaCl) containing 5 % (w/v) BSA. Filters were incubated with purified anti-DevT antiserum in TBS containing 5 % (w/v) BSA, 5 % (v/v) IgG solution and incubated with horseradish peroxidase conjugate (1 : 15 000) (Amersham Biosciences). Immunoreactive bands were detected using the Lumi-Light Western blotting conjugate (1 : 15 000) (Amersham Biosciences). Membranes were blocked in TBS (20 mM Tris pH 7.5, 150 mM NaCl) containing 5 % (w/v) BSA. Filters were incubated with purified anti-DevT antiserum in TBS containing 5 % (w/v) BSA, 5 % (v/v) IgG solution and incubated with horseradish peroxidase conjugate (1 : 15 000) (Amersham Biosciences). Immunoreactive bands were detected using the Lumi-Light Western blotting conjugate (1 : 15 000) (Amersham Biosciences). Membranes were blocked in TBS (20 mM Tris pH 7.5, 150 mM NaCl) containing 5 % (w/v) BSA. Filters were incubated with purified anti-DevT antiserum in TBS containing 5 % (w/v) BSA, 5 % (v/v) IgG solution and incubated with horseradish peroxidase conjugate (1 : 15 000) (Amersham Biosciences). Immunoreactive bands were detected using the Lumi-Light Western blotting conjugate (1 : 15 000) (Amersham Biosciences). Membranes were blocked in TBS (20 mM Tris pH 7.5, 150 mM NaCl) containing 5 % (w/v) BSA. Filters were incubated with purified anti-DevT antiserum in TBS containing 5 % (w/v) BSA, 5 % (v/v) IgG solution and incubated with horseradish peroxidase conjugate (1 : 15 000) (Amersham Biosciences). Immunoreactive bands were detected using the Lumi-Light Western blotting conjugate (1 : 15 000) (Amersham Biosciences).
by loss of fluorescence after 24 h of nitrogen starvation (data not shown). The ultrastructure of filaments from the mutant strain deprived of combined nitrogen for 48 h revealed morphologically mature heterocysts with no conspicuous differences from the wild-type heterocysts (Fig. 1a, b). Both layers of the envelope (glycolipid and polysaccharide layers) were present and the inner membranes were rearranged to form the honeycomb membrane at the poles adjacent to the neighbouring vegetative cells. Both the typical narrow septum between the two cell types and the so-called polar channels are present in the mutant. Therefore, mutant α2 is not impaired in morphological maturation of the heterocysts.

Heterocyst-specific oxidation of DAB, indicating the maturation of the heterocyst protoplast, and reduction of TTC, a sign of heterocyst micro-oxic conditions, were analysed (Fig. 1c). The wild-type strain showed a clear polar DAB staining, indicative of high concentrations of respiratory enzymes located at the honeycomb membranes. By contrast, mutant M7 (a transposon mutant defective in devA: Maldener et al., 1994), used here as control strain, and mutant α2 lacked the polar staining but showed a background of homogeneously precipitated DAB, and were thus considered as DAB⁺, which agrees with previous observations (Ernst et al., 1992). Under our TTC staining conditions, the indicator was reduced rapidly in the heterocysts of the wild-type, resulting in clear formazan crystals, while none were formed in the vegetative cells. The rapid TTC reduction in heterocysts is due to the absence of O₂-evolving photosystem II, the presence of the envelope glycolipid-layer that acts as an O₂ barrier and the increased respiration, leading to a reduced amount of O₂ that otherwise may compete with TTC as an electron acceptor. TTC reduction was not observed in mutant M7, probably due to the absence of the laminated layer (Fiedler et al., 1998). Formazan crystals were hardly detected in heterocysts of mutant α2 and when they appeared, they were significantly smaller than those present in the wild-type heterocysts (Fig. 1c). To investigate whether respiratory activity is high in the mutants following heterocyst differentiation, oxygen consumption of whole filaments was measured after induction of heterocyst differentiation (Table 1). While the wild-type showed a clear increase in respiration during diazotrophic growth, mutants α2 and M7 had comparable low activity after being incubated for 48 h without a source of combined nitrogen. This is in line with the results of DAB and TTC staining, which show, respectively, that the mutants are impaired in respiration and creation of a micro-oxic environment. Since the envelope seems to be fully developed, the inability to provide a micro-oxic environment may be due to insufficient reduction of oxygen by respiration.

Cells of mutant α2 were assayed for their ability to develop nitrogenase activity after nitrogen step-down. Under

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**Table 1. O₂ respiration in Anabaena wild-type, α2 and M7 mutants**

The wild-type strain was grown in the presence of nitrate until the culture reached an OD₆₀₀ of 0.5–0.6; then cells were collected and assayed as described in Methods. In addition, cultures of wild-type and mutant strains were grown with nitrate to similar optical densities (~0.5), then harvested, washed with medium lacking any source of combined nitrogen and further incubated in medium lacking any source of combined nitrogen for 48 h prior to carrying out the measurements.

<table>
<thead>
<tr>
<th>Anabaena strain</th>
<th>O₂ consumption [nmol O₂ (mg Chl a h⁻¹)⁻¹]*</th>
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<tbody>
<tr>
<td>WT (NO₃⁻)</td>
<td>12.15±1.05</td>
</tr>
<tr>
<td>WT (N₂)</td>
<td>50.94±4.74</td>
</tr>
<tr>
<td>α2 (N₂)</td>
<td>19.54±3.48</td>
</tr>
<tr>
<td>M7 (N₂)</td>
<td>19.97±0.97</td>
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</tbody>
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*Values are means±SD of two independent experiments. Chl a, chlorophyll a.
aerobic conditions, filaments of mutant $a2$ showed no reduction of acetylene. However, after incubation under anaerobic conditions (argon, DCMU treatment) they developed about 45% of the activity of the wild-type under the same conditions. These data were similar to those reported by Ernst et al. (1992).

**Recovery and characterization of the mutated genomic region of mutant $a2$**

Plasmid pRL1630 is the circularized _Clal_ fragment recovered from mutant $a2$, bearing the transposon Tn5-1065 flanked by _Anabaena_ DNA; it was obtained from C. Peter Wolk, Michigan State University, East Lansing, USA. The gene where the transposon had inserted was identified by using primers complementary to the ends of the transposon for sequencing. A 9 bp direct repeat (5'-CCGCCAGTG-3') was created upon transposition into the gene. After screening of a λ library of _Anabaena_ chromosomal DNA (Black & Wolk, 1994) with pRL1630 as a probe, the wild-type copy of the gene was cloned on a 7753 bp _Clal_ fragment into vector pBlueScript SK +, resulting in plasmid pLM71. Further subcloning and sequencing revealed an ORF of 957 bp that we named _devT_ (GenBank accession no. CAB57788). _devT_ would encode a protein with molecular mass of 35.33 kDa (319 residues), isoelectric point 5.48) assuming the first ATG would encode a protein with molecular mass of 35.33 kDa (319 residues, isoelectric point 5.48) assuming the first ATG to be the start codon. Thirty base pairs downstream from that codon, a second putative start codon is present. Putative ribosome-binding sites are located upstream from each of those sites. A sequence similar to serW from _E. coli_ encoding tRNA$^{\text{Ser}}$ is found 80 bp downstream of _devT_. In the annotated genome of _Anabaena_ PCC 7120 (Kaneko et al., 2001), _devT_ corresponds to gene _alr4674_. In silico analysis of the _devT_ chromosomal region does not show additional ORFs that could form a transcription unit with _devT_, except for the tRNA-Ser downstream of _devT_ (Fig. 2a).

Reconstruction of the _devT_ mutant in the wild-type of _Anabaena_ was achieved by site-directed mutagenesis of _devT_. The neomycin-resistance cassette C.K3 (Elhai & Wolk, 1988a) was cloned in the opposite orientation with respect to _devT_ transcription into the unique _NdeI_ site of _devT_ (Fig. 2a). After transfer into _Anabaena_ cells, double recombinants were selected by a positive selection method based on the use of the conditionally lethal gene sacB (Cai & Wolk, 1990). Disruption of the gene was confirmed by Southern blot analysis and PCR (data not shown). The recombinant strain DRpMA54 showed the same phenotype as the original mutant _a2_ and was not able to grow on N$_2$, although heterocysts were formed upon nitrogen starvation. The ultrastructure observed by transmission electron microscopy was similar to that of mutant _a2_ and the wild-type (data not shown). pDU1-based shuttle vector pMA40, harbouring the entire ORF of _devT_ including flanking regions, was transferred into strains DRpMA54 and _a2_, resulting in complemented strains that had recovered the ability to grow on N$_2$ as sole nitrogen source (shown for _a2C_ in Fig. 2b).

**_DevT_ is a Ser/Thr phosphatase conserved in the photosynthetic lineage**

The deduced amino acid sequence of _DevT_ shows homology to the family of phospho-Ser/Thr phosphatases, which belongs to the superfamily of phosphoesterases (Koonin, 1994). A region of 217 amino acids (residues 21–237) shows high similarity to the catalytic domain of PPP-type Ser/Thr protein phosphatases and contains the three conserved motifs of PPP-type phosphatases (see Supplementary Fig. S1, available with the online version of this paper) (Barton et al., 1994; Shi et al., 1998; Shi, 2009). _DevT_ contains the conserved residues involved in metal binding (motifs I and II) and the catalytic histidine residue (motif III), respectively (Barford, 1996). While motif I of _DevT_ matches exactly the consensus sequence, motif II is less conserved in _DevT_. However, it contains the Gly and Asp residues of the consensus sequence GD(LYF)V(DA)RG, proposed by Barton et al. (1994). The RG residues in this motif do not as highly conserved as the GD residue pair. Amino acid residues between the three motifs do not show much conservation among different phosphoesterases. The previously identified PPP-type phosphatase PrpA from _Anabaena_ shows low similarity to _DevT_.

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**Fig. 2.** The _devT_ gene cluster and Fox phenotype of _devT_ mutant strains. (a) Genomic organization of the chromosomal region containing the _devT_ gene (_alr4674_) and the two strategies to inactivate _devT_: via random insertion of transposon Tn5-1065 and via insertion of the C.K3 resistance cassette (mutants _a2_ and DRpMA54, respectively). (b) Growth on Allen & Arnon agar plates containing 5 mM NaNO$_3$ (+N) or lacking combined nitrogen (–N) of wild-type (WT), _devT_ (_a2_ and DRpMA54) and _devT_-complemented (_a2C_) strains.
Biochemical characterization of DevT

The devT gene was cloned into the pQE70 vector (yielding plasmid pIM88) to produce a recombinant DevT protein with its C-terminus fused to six histidine residues. After overproduction (see Methods), most of the protein was in the insoluble pellet of the crude cell extracts. Nevertheless, the soluble fraction contained enough DevT protein to be purified on a Ni²⁺-NTA column (Fig. 3a). To analyse the native molecular mass of DevT-H₆, size-exclusion chromatography on a Superdex G200 10/300 GL column was performed. The purified fusion protein eluted in one peak corresponding to a molecular mass of about 36 kDa, which would correspond to monomeric DevT protein in solution (Fig. 3b). N-terminal sequencing of the DevT purified protein gave an amino acid sequence exactly matching that of the predicted protein, if the assumed first start codon (ATG) is used in E. coli.

Given that the deduced amino acid sequence contains the metal-binding and active-site residues of PPP-type phosphatases, we analysed whether DevT shows phosphorylase activity in vitro using the commonly used chromogenic substrate pNPP. The purified DevT-H₆ protein did indeed show phosphatase activity with pNPP as substrate, but only in the presence of Mn²⁺ (Fig. 4a). The estimated Kₘ for Mn²⁺ was 23.6 μM (estimation by Hanes–Woolf plot of the direct plot shown in Fig. 4b) and the apparent kinetic constants of the DevT activity towards pNPP corresponded to a Kₘ of 2.5 mM and a Vₘₐₓ of 0.8 μmol pNPP min⁻¹ mg⁻¹ (Hanes–Woolf plot, not shown). However, DevT was not able to remove the phosphate from a peptide containing phosphothreonine RRA(pT)VA or phosphoserine RRA(pS)VA (Promega). Furthermore, DevT was not able to dephosphorylate phosphorylated PII protein from Synechococcus elongatus in vitro nor 32P-casein (not shown).

DevT activity was maximal at 55 °C and at a pH of 8 (data not shown). The ability of phosphatase inhibitors to influence DevT activity was analysed (Fig. 4c). Addition of the competitive inhibitor inorganic phosphate and the metal chelator EDTA resulted in complete inhibition, but vanadate, NaF and tartrate also decreased DevT activity.

Because it is known that some protein phosphatases are redox regulated (Rusnak & Reiter, 2000), we measured pNPP hydrolysis after incubation of purified DevT protein under different redox conditions. Oxidation of DevT by
either incubation in the presence of increasing concentrations of H₂O₂ or prolonged exposure to H₂O₂ resulted in a significant decrease of activity (Fig. 4d, e), with a half-maximal inhibitory concentration (IC₅₀) of 250 μM H₂O₂. However, subsequent incubation with the reducing agent DTT restored the pNPP dephosphorylation activity (Fig. 4f).

The devT gene is constitutively expressed but DevT protein accumulates only during diazotrophic growth

Since mutant a2 was impaired in diazotrophic growth, we investigated whether devT expression is differentially regulated upon nitrogen deprivation and whether it is affected by the two master regulators of heterocyst differentiation, NtcA and HetR. Northern blot analysis was performed using total RNA isolated from filaments of the wild-type Anabaena strain, ntcA mutant CSE2 (Frias et al., 1994) and hetR mutant DR884a (Black et al., 1993) at different time intervals after nitrogen step-down. No substantial difference in the abundance of the mRNA specifically hybridizing to the devT probe could be observed during nitrogen step-down. No substantial difference in the abundance of the mRNA specifically hybridizing to the devT probe could be observed during nitrogen step-down in wild-type cells (Fig. 5a, WT lanes). Correspondingly, neither upregulation nor downregulation of devT was observed by RT-PCR in wild-type cells starved for nitrogen for up to 48 h (Fig. 5b). ntcA and hetR mutants showed similar and a moderate increase of devT expression, respectively, in comparison with wild-type (Fig. 5a, NtcA⁻ and HetR⁻ lanes).

Immunoblot analyses were carried out to analyse DevT protein levels in wild-type Anabaena cells deprived of combined nitrogen. The polyclonal antisera prepared against DevT detected a protein in wild-type extracts that was absent from mutant a2 (data not shown). The estimated mass of this protein, roughly 36 kDa, is in agreement with the value predicted from the deduced amino acid sequence of DevT. Total soluble protein was isolated from filaments of cells shifted from ammonia-supplemented to combined nitrogen-free medium, with samples collected at different time points (Fig. 6a). DevT was detected in extracts from wild-type whole filaments deprived of nitrogen for 48 h, but not at 9, 24 or 34 h. When proteins from isolated heterocysts were separated by SDS-PAGE and blotted, DevT was first noticeable after 24 h and increased strongly during the next 24 h (Fig. 6b).

Heterocyst-specific localization of DevT

Localization of DevT in the diazotrophic filaments of Anabaena wild-type strain was examined by using the green fluorescent protein (GFP) as a reporter. The devT–gfp fusion was inserted into a neutral site (nucA–nuIA region) located in the α-megaplasmid, essentially as
described by Olmedo-Verd et al. (2006). The selected clone (strain WT-333) bore a copy of the devT–gfp construct, integrated in the α-megaplasmid, and retained an intact copy of the wild-type devT region, as verified by PCR with specific oligonucleotides (data not shown). DevT–GFP accumulated in fully developed heterocysts, which showed a decreased autofluorescence of photosynthetic pigments. Faint green fluorescence in heterocysts from the reporter strain could be observed 28 h after nitrogen step-down (Fig. 7). Subsequently, heterocysts showed a progressive increase of GFP fluorescence, with the highest level in heterocysts of steady-state N2-fixing cultures (Fig. 7). These results suggest that the accumulation of DevT takes place specifically in mature, presumably older, heterocysts, at least up to a certain heterocyst age.

**DISCUSSION**

In this study, we characterized devT (alr4674) from *Anabaena*, which encodes a PPP-family phosphatase involved in heterocyst function. Orthologues of DevT were found in the available genomes of cyanobacteria and certain genomes of plants and the green alga *Chlamydomonas*. Cyanobacterial DevT sequences included heterocyst-forming (as *Anabaena variabilis* ATCC 29413), nitrogen-fixing non-heterocystous (as *Trichodesmium erythraeum*) strains and filamentous non-fixing (as *Lyngbya* sp. PCC 8106) as well as unicellular strains (as *Synechocystis* sp. PCC 6803), which do not fix N2. The conservation of DevT in strains with different cell structure and metabolism suggests different functions for DevT in these organisms. Since close DevT homologues are only present in organisms that perform oxygenic photosynthesis, its role could also be related to redox control and oxygen stress.

DevT shows a manganese-dependent phosphatase activity, and the effect of various inhibitors of different families of protein phosphatases together with sequence-based
similarities of the catalytic domain are consistent with DevT being a member of the PP1 family. pNPP hydrolysis was observed using recombinant DevT, but the protein was not able to remove the phosphate from peptide substrates nor from casein. Several known serine/threonine phosphatases, such as PP2A, PP2B and PP2C, dephosphorylate those substrates. However, the artificial phospho-peptides used in this study are poor substrates for PP1 phosphatases given their more stringent structural requirements, and this could also be the case for the lack of activity of DevT-H6. Also, phosphorylated PII protein was not a substrate for DevT-H6. Further research should clarify whether DevT can act on targets containing phospho-Thr or phospho-Ser. Due to conservation in the primary sequences of phosphoesterases it cannot be excluded that DevT acts on non-proteinaceous substrates, such as cyclic-nucleotide phosphates, polynucleotides, etc. (Koonin, 1994).

In the last two decades of heterocyst research, the approach of reverse genetics by transposon mutagenesis has proved to be a straightforward tool for identifying new genes involved in this prokaryotic differentiation process (e.g. Ernst et al., 1992; Fan et al., 2005). Transposon mutant a2 (a devT mutant), identified previously by its inability to grow on N2 as sole nitrogen source, has been analysed in depth here. The ultrastructure of the non-functional heterocysts of the mutant showed no differences from that of the wild-type, in contrast to the first description by (Ernst et al., 1992). Hence the devT mutant could be altered in regulatory or metabolic processes. In this respect, DAB and TTC staining as well as respiration measurements suggest that the drop in O2 concentration that takes place during heterocyst formation does not occur efficiently in the devT mutant, leading to a defect in nitrogenase activity that results in the inability to grow diazotrophically. It should be noted that mutant a2 showed a severe decrease of nitrogenase protein levels in the presence of air compared to anaerobic conditions, the latter conditions partially restoring nitrogenase activity and levels (Ernst et al., 1992).

DevT protein was shown to be upregulated and accumulated only during diazotrophic growth, specifically in heterocysts. While the mRNA of devT stayed at a similar low level during nitrogen step-down, low levels of DevT protein were initially detected at a time when heterocysts became mature and functional, about 24 h after step-down. After 48 h, devT mRNA levels remained at a

Fig. 7. Specific accumulation of the DevT-GFP translational fusion in heterocysts. Accumulation of DevT in heterocysts after nitrogen step-down, as reported by the GFP-based fluorescence at indicated time points of *Anabaena* bearing a devT–gfp fusion integrated into the α-megaplasmid. Arrows indicate the position of heterocysts. Bright-field (top), red autofluorescence (middle) and GFP fluorescence (bottom) images are shown.
constant low level whereas the protein significantly accumulated. Indeed, transcription of the \textit{devT} gene was constitutive under tested conditions, independent of NtcA, while HetR exhibited a certain negative effect. Taken together, our data suggest that DevT might be post-transcriptionally regulated, and there might be a mechanism that controls stability of the DevT protein. DevT protein could be easily degraded in vegetative cells and immature heterocysts (proheterocysts), but – by an unknown mechanism – might be stabilized in mature heterocysts. Whether HetR constitutes an additional point of transcriptional regulation of \textit{devT} directly by binding to regulatory DNA elements or indirectly requires further investigation.

Considering the phenotypic features of the \textit{z2} mutant and the late accumulation of DevT during heterocyst formation, DevT seems to be a factor involved in late steps of heterocyst maturation (approx. 24 h after initiation of heterocyst formation), presumably transmitting signals, via dephosphorylation, that are essential to achieve functionality in mature heterocysts. Our data clearly indicate that the phenotype of the \textit{devT} mutant is related to oxygen stress. In order to understand the role of DevT, the signals transmitted by DevT as well as its putative target or targets need to be addressed.

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