

Two strictly polyphosphate-dependent gluco(manno)kinases from diazotrophic *Cyanobacteria* with potential to phosphorylate hexoses from polyphosphates

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Abstract The single-copy genes encoding putative polyphosphate–glucose phosphotransferases (PPGK, EC 2.7.1.63) from two nitrogen-fixing *Cyanobacteria*, *Nostoc* sp. PCC7120 and *Nostoc punctiforme* PCC73102, were cloned and functionally characterized. In contrast to their actinobacterial counterparts, the cyanobacterial PPGKs have shown the ability to phosphorylate glucose using strictly inorganic polyphosphates (polyP) as phosphoryl donors. This has proven to be an economically attractive reagent in contrast to the more costly ATP. Cyanobacterial PPGKs had a higher affinity for medium–long-sized polyP (greater than ten phosphoryl residues). Thus, longer polyP resulted in higher catalytic efficiency. Also in contrast to most their homologs in *Actinobacteria*, both cyanobacterial PPGKs exhibited a modest but significant polyP-mannokinase activity as well. Specific activities were in the range of 180–230 and 2–3 $\mu\text{mol min}^{-1} \text{mg}^{-1}$ with glucose and mannose as substrates, respectively. No polyP-fructokinase activity was detected. Cyanobacterial PPGKs required a divalent metal cofactor and exhibited alkaline pH optima (approx. 9.0) and a remarkable thermostability (optimum temperature, 45 °C). The preference for Mg^{2+} was noted with an affinity constant of 1.3 mM. Both recombinant PPGKs are homodimers with a subunit molecular mass of ca. 27 kDa. Based on database searches and experimental data from Southern blots and

activity assays, closely related PPGK homologs appear to be widespread among unicellular and filamentous mostly nitrogen-fixing *Cyanobacteria*. Overall, these findings indicate that polyP may be metabolized in these photosynthetic prokaryotes to yield glucose (or mannose) 6-phosphate. They also provide evidence for a novel group-specific subfamily of strictly polyP-dependent gluco(manno)kinases with ancestral features and high biotechnological potential, capable of efficiently using polyP as an alternative and cheap source of energy-rich phosphate instead of costly ATP. Finally, these results could shed new light on the evolutionary origin of sugar kinases.

Keywords Gluco(manno)kinase · Hexokinase · *Cyanobacteria* · PolyP-dependent · Polyphosphate · Regeneration of ATP

Introduction

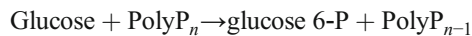
Inorganic polyphosphate (polyP) is a linear polyanion composed of tens to hundreds of phosphoryl residues, all of them being linked by “high-energy” phosphoanhydride bonds. Found in many diverse organisms in nature, polyP has proven to be essential for the growth, response to stresses, and stringencies of cells (Kulaev 1979; Wood and Clark 1988; Kornberg et al. 1999; Rao et al. 2009).

As phosphorylated compounds with a Gibbs free energy of hydrolysis similar to the ATP ($-30.5 \text{ kJ mol}^{-1}$), inorganic pyrophosphate (diphosphate, PPi) and polyP have been suggested to be used in place of ATP in diverse biological processes (Lipmann 1965). An enzyme known to hydrolyze polyP rather than ATP is the polyphosphate glucokinase (PPGK, polyphosphate–glucose phosphotransferase, EC

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2.7.1.63), which catalyzes the phosphorylation of glucose using polyP as a phosphoryl donor to form glucose 6-phosphate as follows:



PPGK was first observed in *Mycobacterium phlei* (Szymona 1957), and later in other Gram-positive bacteria, all of them belonging to the ancient order of *Actinomycetales* (Szymona and Ostrowski 1964; Szymona and Widomski 1974; Szymona and Szymona 1978, 1979; Pepin and Wood 1986; Mukai et al. 2003; Tanaka et al. 2003; Lindner et al. 2010a; Hehuan et al. 2012; Koide et al. 2013). However, no PPGK has been described in other sorts of bacteria, archaea, fungi, algae, plants, or animals to date.

Most actinobacterial PPGKs have been validated as monomers or homodimers with a molecular mass subunit of ca. 30 kDa. A remarkable feature of these enzymes is its dual substrate specificity: PPGK can use both ATP and polyP as donors to phosphorylate glucose to glucose 6-phosphate. Nevertheless, a PPGK from the polyP-accumulating actinobacterium *Microtholunatus phosphovorius* is the only PPGK enzyme solely dependent on polyP as an energized phosphoryl substrate donor described to date (Tanaka et al. 2003). Concerning this matter, several studies (Hsieh et al. 1993; Phillips et al. 1999) proved that although both enzymatic activities belong to the same protein, the binding site for this protein differs in each phosphate donor substrate. Thus, the enzymes of most phylogenetically ancient species of the *Actinomycetales* order seem to prefer polyP instead of ATP. As a result of this, a higher polyP glucokinase/ATP glucokinase ratio is exhibited. Compared to the rest of glucokinases, PPGKs displayed a wider range of nucleoside 5'-triphosphates (NTP) as phosphoryl donors [guanosine 5'-triphosphate (GTP), uridine 5'-triphosphate (UTP), thymidine 5'-triphosphate (TTP), xanthosine 5'-triphosphate (XTP), cytidine 5'-triphosphate (CTP) and deoxyadenosine 5'-triphosphate (dATP)], whereas ATP glucokinases from more evolved organisms are unable to use polyP and, consequently, only poorly replace GTP for ATP, as is the case of hexokinases (EC 2.7.1.11) from fungi and mammals, which are exclusively dependent on ATP (Rao et al. 2009).

PPGK belongs to the repressor ORF kinase (ROK) superfamily (Pfam PF00480) (Finn et al. 2014), a large group of mostly bacterial proteins which also include other sugar kinases and transcriptional repressors, the latter with an extra h- α -h DNA binding domain. Owing to this fact, kinase enzymes within this group (bacterial gluco-, fructo-, and mannokinases; eukaryotic hexokinases; and ADP glucokinases) reveal a significant grade of structural relationship.

PolyP should play important roles in the overcoming of nutrient and heavy metal stresses by *Cyanobacteria*, a group of Gram-negative oxygenic photoautotrophic prokaryotes

which are among the most successful and oldest forms of life (Schopf 2002) and have gained a lot of attention in recent years because of their potential applications in biotechnology (Abed et al. 2009). Accumulation of polyP granules has been described under various culture conditions in both unicellular (Lawry and Jensen 1979) and filamentous (Jensen et al. 1982) *Cyanobacteria*. Moreover, induction of genes involved in polyP metabolism by Pi starvation was reported in the unicellular strain *Synechocystis* sp. (Gomez-Garcia et al. 2003). Interestingly, in the diazotrophic filamentous cyanobacterium *Anabaena flos-aquae*, P is stored in different ways depending of the nitrogen source used. Under dinitrogen-fixing conditions, P is stored as sugar P, whereas with nitrate as the combined N source, it is stored as polyP (Thompson et al. 1994). However, the functional relationships between polyP metabolism and dinitrogen fixation in cyanobacteria have not yet been elucidated.

Here, we report the first polyP gluco(manno)kinases isolated and biochemically characterized from *Cyanobacteria*. The single-copy *ppgK* genes of the filamentous nitrogen-fixing strains *Nostoc* sp. PCC7120 and *Nostoc punctiforme* PCC73102 were cloned and overexpressed in *Escherichia coli*, and the corresponding recombinant proteins, hereafter referred as NsPPGK and NpPPGK, respectively, were purified and characterized. As shown, these enzymes are smaller proteins and exhibit some novel biochemical features compared to the previously described PPGKs. Additionally, a survey of homologous closely related PPGKs has been carried out in a wide range of diazotrophic *Cyanobacteria* by several techniques including Southern blots, activity assays, and bioinformatic analyses. Lastly, this study could also offer new evidence towards the matter of hexokinase evolution. Overall, the obtained results provide indications for cyanobacterial PPGKs representing a taxonomic group-specific new subfamily of strictly polyP-dependent gluco(manno)kinases with high biotechnological potential.

Materials and methods

Reagents and PolyP preparation

Restriction endonucleases and T4 DNA ligase were purchased from Takara Bio Inc (Shiga, Japan). Accuzyme™ proofreading DNA polymerase and the gel extraction kit were obtained from Bioline Inc. (MA, USA). Primers were synthesized by Integrated DNA Technologies (Leuven, Belgium). Sodium polyphosphates P₁i, tripolyphosphate (P₃), cyclic P₃ (trimetaphosphate, P_{3c}), and tetrapolyphosphate (P₄); a polyphosphate mix with an average chain length of 13–18 phosphoryl residues (P_{13–18}) and water-insoluble Maddrell salt (a mixture of crystalline long-chain polyphosphates of very high molecular mass); NTPs; dATP; and hexoses (D-

glucose, D-mannose, D-fructose) were purchased from Sigma Chemical Co. (St. Louis, MO, USA). Purchased substrates (polyPs and hexoses) were analytical grade reagents, except the Maddrell salt which was of practical grade. P₆₀ and P₁₅₀ (polyphosphate mixes purified by polyacrylamide gel electrophoresis; average chain lengths 60 and 150 phosphoryl residues, respectively) were kindly provided by Dr. Toshikazu Shiba (RegeneTiss Co., Japan). Very long-chain polyPs with chain lengths of up to approximately 800 phosphoryl residues (P_{LC}) were obtained by fractionation of solubilized Maddrell salt, prepared as described by Van Wazer (1958) on a 2 % (w/v) polyacrylamide/0.8 % (w/v) agarose gel. When necessary, crystalline polyP was washed twice with 70 % (v/v) ethanol, dried overnight in a vacuum desiccator, and resuspended in distilled water. Otherwise stated, the polyP concentration is expressed in terms of polymer, assuming average chain lengths of 3, 4, 15, 60, 150, and 300 phosphoryl residues for P₃, P₄, P_{13–18}, P₆₀, P₁₅₀, and P_{LC}, respectively. All other chemicals were of analytical grade.

Analytical polyacrylamide gel electrophoresis of polyP

Polyacrylamide slab gels (total acrylamide, 30 %, w/v; 70×85 mm; 1 mm thick) were prepared which contained a 19.2:0.8 ratio of acrylamide to bisacrylamide. The gel was pre-electrophoresed at 100 V for 3 h to remove contaminating ions. The polyP samples were mixed at a ratio of 1:6 with loading buffer [100 mM Tris–borate buffer (pH 8.3), 30 % (v/v) glycerol, and 0.25 % (w/v) bromophenol blue]. Gels were run at 50 mA in Tris–borate–EDTA (TBE) as electrophoresis buffer. Once electrophoresed, polyPs were fixed and stained with 0.05 % (w/v) toluidine blue O, 25 % (v/v) methanol, and 1 % (v/v) glycerol in water, followed by destaining in an aqueous mixture containing 25 % (v/v) methanol and 5 % (v/v) glycerol. As a result, the polyP stained dark blue against the colorless or light blue background.

Bacterial strains and culture conditions

The cyanobacterial strains used in this work were obtained as axenic cultures from various microbial culture collections of reference for *Cyanobacteria* (American Type Culture Collection (ATCC), Manassas, VA, USA; Pasteur Culture Collection (PCC), Paris, France; Culture Collection of Algae, University of Texas (UTEX), Austin, TX, USA; the CICC Artuja Biological Cultures Service, Instituto de Bioquímica Vegetal y Fotosíntesis, Seville, Spain). The strains were photoautotrophically grown in BG11 liquid medium without combined nitrogen source unless otherwise stated (Rippka et al. 1979) and are described in Online Resource Table S1. Cultures (referred to as bubbled cultures) were supplemented with 10 mM NaHCO₃ and bubbled with a mixture of CO₂ and air (2 %, v/v), under continuous

fluorescent white light (75 μE m⁻² s⁻¹). The absence of heterotrophic bacterial contamination was assessed by counts on Luria–Bertani (LB) agar plates incubated in the dark.

Preparation of cyanobacterial cell-free extracts

Cyanobacterial cells were harvested by centrifugation and resuspended in 100 mM Tris–HCl (pH 9.0) buffer supplemented with 5 mM MgCl₂, 5 mM DTT, 0.1 mM PMSF, and a 1:1000 dilution of a protein inhibitor cocktail for use with bacterial cell extracts (P8465; Sigma-Aldrich, USA), at a ratio of 0.2 g (cell wet wt)/ml. Then, cells were ultrasonically disrupted at 0–4 °C. The cell homogenate was centrifuged at 15,000×g at 4 °C for 20 min, and the resultant clear supernatant (cell-free extract) was used for enzymatic assays.

DNA methodology

Total DNA was isolated by the following procedure: 50 ml of cyanobacterial cultures in the mid-log phase of growth was harvested and resuspended in a final volume of 400 μl in a microcentrifuge tube with 10 mM Tris–HCl (pH 7.5) buffer with 0.1 mM EDTA. Then, 150 μl of sterile glass beads (0.2 μm diameter), 20 μl of 10 % (w/v) sodium dodecyl sulfate (SDS), and 450 μl of phenol–chloroform–isoamyl alcohol mixture (25:24:1 v/v) were added. The mixture was subjected to six cycles of 1-min vigorous vortexing followed by 1-min cooling on ice. The resulting suspension was centrifuged at 15,000×g for 10 min; then, the clear supernatant solution was transferred to a new microcentrifuge tube, and DNA was finally ethanol precipitated.

Southern blotting

DNA samples isolated from a number of strains representative of the different taxonomic groups of cyanobacteria were digested with appropriate restriction enzymes and loaded onto agarose gels; then, Southern analysis was performed (Ausubel et al. 1992) using GeneScreen Plus membranes (DuPont, USA). DNA probes utilized in the hybridizations (full coding *ppgK* fragments) were obtained by PCR and were then labeled with [α -³²P]-dCTP using the Ready-To-Go® DNA labeling kit (GE Healthcare). Nucleic acid hybridization was carried out at 55 °C with gentle shaking. Films were exposed for 4 days and developed using a Cyclone® Storage Phosphor System (Packard, USA).

Construction of recombinant plasmids and gene expression in *E. coli*

The *ppgK* genes from *Nostoc* sp. PCC7120 and *N. punctiforme* PCC73102 were PCR amplified using specific primers (Online Resource Table S2) and genomic DNA as a

template. The unique DNA fragments of ca. 0.72 kb obtained in both cases were initially cloned into the pGEM-T Easy vector (Invitrogen) for sequencing. These plasmids were then digested with *Bam*HI and *Pst*II, and the DNA fragments carrying the native open reading frames of *ppgK* genes were eventually ligated into pQE-80L vector (Quiagen, Germany). In this way, a His₆ tag of 12 amino acid residues in total (MRGSHHHHHHGS; nominal mass 1420 Da) was added to the N-terminal end of the native proteins. *E. coli* BL21(DE3) cells transformed with the appropriate expression plasmid were cultured at 30 °C in 1 l LB liquid medium supplemented with 100 µg ml⁻¹ ampicillin with vigorous shaking. When OD₆₀₀ reached ca. 0.6, protein expression was induced by adding 1 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) and cultures were then incubated overnight at 20 °C with shaking at 200 rpm.

Purification of recombinant cyanobacterial PPGKs by nickel-nitrilotriacetic acid (Ni-NTA) metal-affinity chromatography

Cells were harvested and resuspended in buffer A (500 mM NaCl, 50 mM Na₂HPO₄, 10 mM imidazole, pH 8.0) and then lysed by sonication at 4 °C. Cell debris were removed by centrifugation at 15,000×g for 15 min. The resultant crude extract was loaded onto a pre-equilibrated HisTrap FF Crude Ni-NTA 1-ml column (GE Healthcare). Subsequently, nontarget proteins were removed by washing the column with buffer B (500 mM NaCl, 50 mM Na₂HPO₄, 50 mM imidazole, pH 8.0) until no more protein elution was observed. Finally, recombinant proteins were eluted by applying a linear gradient with a target concentration of 100 % of buffer C (500 mM NaCl, 50 mM Na₂HPO₄, 500 mM imidazole, pH 8.0). The eluted PPGK proteins were dialyzed three times with 50 mM Tris-HCl (pH 9.0) to remove imidazole and phosphate salts and eventually concentrated by ultrafiltration using Amicon Ultra 3-kDa filters.

Fast protein liquid chromatography gel filtration: estimation of molecular masses

Partially purified His-tagged PPGK preparations, previously concentrated by ultrafiltration, were further purified by fast protein liquid chromatography (FPLC) gel filtration carried out at 4 °C. The concentrated preparations (0.5–1.0 ml volume) were loaded on to a Superdex© 200 PG (GE Healthcare, Sweden) column equilibrated with 150 mM NaCl, 20 mM KCl, 5 mM MgCl₂, and 50 mM Tris-HCl (pH 9.0) buffer at a flow rate of 2 ml min⁻¹ using an ÄKTA-FPLC system (GE Healthcare, Sweden). The molecular masses (*M_m*) of oligomeric PPGK proteins were determined using the calibration plot derived from the elution volumes of a series of protein standards including thyroglobulin (Thy, 669 kDa), ferritin

(Fer, 443 kDa), β-amylase (β-Amy, 200 kDa), alcohol dehydrogenase (ADH, 150 kDa), bovine serum albumin (BSA, 66 kDa), carbonic anhydrase (CA, 29 kDa), and cytochrome c (Cyt.c, 12.4 kDa). Subunit molecular masses were determined by denaturing discontinuous SDS-polyacrylamide gel electrophoresis (PAGE) following the method of Laemmli (1970) using 12 % (w/v) separating and 4 % (w/v) stacking polyacrylamide gels. Protein bands were stained with Coomassie Brilliant Blue R-250. Apparent *M_m* of monomers under denaturing PAGE was calculated using standard proteins. Absolute *M_m* values of purified recombinant PPGKs were confirmed by matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometry (see below). These purified fractions were used for the in vitro kinetics assays and biochemical characterization.

Peptide mass fingerprinting and validation of PPGK proteins by MALDI-TOF mass spectrometry

Protein samples corresponding to high-purity cyanobacterial PPGKs were derived from SDS-PAGE. Proteins were digested with trypsin, and the resulting peptides were extracted and then loaded onto a suitable MALDI matrix and eventually processed using a MALDI-TOF mass spectrometer (AutoFlex, Bruker-Daltonics, Proteomics Service of the Instituto de Bioquímica Vegetal y Fotosíntesis, CSIC and University of Seville) which generated peptide mass spectra in the mass range 0.8–2.5 kDa. Mascot Matrix Science database was used to analyze the peak lists for protein identification (Koenig et al. 2008).

Determination of enzymatic activities

Unless otherwise stated, sugar kinase enzymatic activities were determined at 40 °C and pH 9.0, using P_{13–18} as a phosphoryl donor substrate. The polyP glucokinase activity was assayed spectrophotometrically by monitoring the production of NADPH at 340 nm using a glucose 6-phosphate dehydrogenase-coupled reaction. The assay mixture (1 ml) contained of 100 mM Tris-HCl buffer (pH 9.0), 5 mM MgCl₂, 5 mM glucose, 1.11 mM polyP, 5 mM nicotinamide adenine dinucleotide phosphate (NADP⁺), and 0.5 U of yeast glucose 6-phosphate dehydrogenase (Sigma Chemical Co., USA). The reaction was started by the addition 0.5–1.5 µg of purified PPGK or 10–20 µl of cell-free extracts. Concentrations of polyphosphate substrates were calculated as polymers, considering mean chain lengths of 15, 60, and 300 phosphate residues for P_{13–18}, P₆₀, and P_{LC}, respectively. NTPs were used at 2 mM concentration when assayed as alternative phosphoryl donor substrates instead of polyP. To determine the dependence on pH, 1.0 µg of purified enzyme was incubated as described above in the following buffers at

100 mM concentration: 2-morpholinoethanesulfonic acid (MES) (pH 5.5–7.0), 3-(N-morpholino)-1-propane sulfonic acid (MOPS) (pH 7.0–8.0), Tris (pH 8.0–9.0), *N*-cyclohexyl-2-aminoethanesulfonic acid (CHES) (pH 9.0–10.0), and 3-[cyclohexylamino]-1-propane sulfonic acid (CAPS) (10.0–10.5). When measuring enzymatic activity in cell-free extracts or when the effects of pH, temperature, divalent metal ions, inhibitors, and other factors on glucokinase activity were examined, the assay was discontinued and NADP⁺ and glucose 6-phosphate dehydrogenase were omitted from the assay mixture. The reaction was finished by heating the test tube at 95 °C for 5 min. Then, the assay followed as described above by adding 5 mM NADP⁺ and 0.5 U of glucose 6-phosphate dehydrogenase. The polyP mannokinase activity was assayed in a similar way, but glucose was replaced by 50 mM mannose and 0.5 U of mannose 6-phosphate isomerase (from *E. coli*; Sigma Chemical Co., USA). Finally, for fructokinase activity determinations, 50 mM fructose and 0.5 U of yeast glucose 6-phosphate isomerase (Sigma Chemical Co., USA) were added in substitution of glucose. Kinetic parameters (K_m and k_{cat}) were determined from initial velocity data that were fitted by the nonlinear regression software Anemona.xlt (Hernandez and Ruiz 1998). One unit (U) of PPGK corresponds to 1 μ mol of phosphorylated product per minute at 30 °C. Protein concentration was determined using the Bradford method (Bradford 1976) with ovalbumin as a standard.

Computer-aided analysis

Amino acid sequence homology among the PPGK sequences was analyzed online using Basic Local Alignment Search Tool (BLAST) searches (Altschul et al. 1990) against the public databases GenBank (Benson et al. 2013), DOE Joint Genome Institute (JGI) (Nordberg et al. 2014), and InterPro (Hunter et al. 2011). The amino acid sequences of putative PPGK orthologs from diverse bacterial strains (Online Resource Table S3) were aligned, and phylogenetic trees were constructed with the evolutionary distances (neighbor joining), maximum parsimony, and maximum likelihood methods using the SeaView v5.2 software (Gouy et al. 2010).

Nucleotide sequence accession numbers

The nucleotide sequences of the gene constructs reported in this paper have been deposited in the GenBank/EMBL/DBJ nucleotide sequence databases under accession numbers HG764586 (*ppgK* of *Nostoc* sp. PCC7120) and HG764587 (*ppgK* of *N. punctiforme* PCC73102), respectively.

Results

all1371 and Npun_R1878 genes encode functional polyP-dependent glucokinases

BLAST sequence similarity searches in cyanobacterial genomes (CyanoBase, Kazusa DNA Research Institute) (Fujisawa et al. 2014) identified two ORFs, *all1371* and *Npun_R1878* of the diazotrophic filamentous strains *Nostoc* sp. PCC7120 and *N. punctiforme* PCC73102, respectively, with high homology to the *ppgK* gene from *Mycobacterium tuberculosis* H₃₇Rv (Hsieh et al. 1996a). The corresponding predicted proteins, thereafter named NsPPGK and NpPPGK, shared 32 and 29 % identity with their mycobacterial homolog and 91 % sequence identity to each other. In addition, each of the genomes of *Nostoc* sp. PCC7120 and *N. punctiforme* PCC73102 possessed a gene encoding a putative glucokinase, *alr2973* and *Npun_R5075*. They respectively showed 27 and 14 % sequence identity at the protein level with their corresponding PPGK homolog. Even though both sequences of putative *ppgK* genes were available, *Npun_R1878* was wrongly annotated as a transcriptional regulator/sugar kinase (ROK family protein) instead of a PPGK-encoding gene. The predicted NsPPGK and NpPPGK polypeptides have 239 (nominal mass 25,919 Da) and 238 (nominal mass 25,816 Da) amino acid residues, respectively. They are smaller than their actinobacterial homologs (of 260–280 residues) and exhibit in their primary structures the seven regions with structural motifs conserved among the bacterial PolyP/ATP-dependent PPGKs (Mukai et al. 2003), as revealed by protein sequence alignments carried out using the ClustalW program (Larkin et al. 2007). Interestingly, when other putative cyanobacterial PPGK sequences were used in the alignment, a high level of conservation was found within them, while when cyanobacterial PPGKs are compared to their actinobacterial polyP/ATP-dependent homologs, motifs reported to be involved in phosphoryl donor and polyphosphate substrate binding (phosphate-1 and phosphate-2, connect-1) and the glucose-binding motif are more clearly conserved (Online Resource Fig. S1). Thus, the finding of putative *ppgK* genes led us to investigate whether glucose 6-phosphate synthesis in *Cyanobacteria* could take place enzymatically through a similar way to that previously described in *M. tuberculosis* along with other *Actinobacteria*. To characterize NsPPGK and NpPPGK, their respective putative genes were obtained from genomic DNA by PCR amplification which yielded a single product with the expected size of 0.72 kb in both cases (Fig. 1a). They were lastly cloned into the pQE-80L expression vector and overexpressed in *E. coli* (BL21). Protein expression was induced in early log-phase cultures by addition of IPTG. The heterologous overexpression of cyanobacterial *ppgK* genes conferred high PPGK activity to *E. coli* cells. Thus, crude extracts from induced *E. coli* cells overproducing NsPPGK or

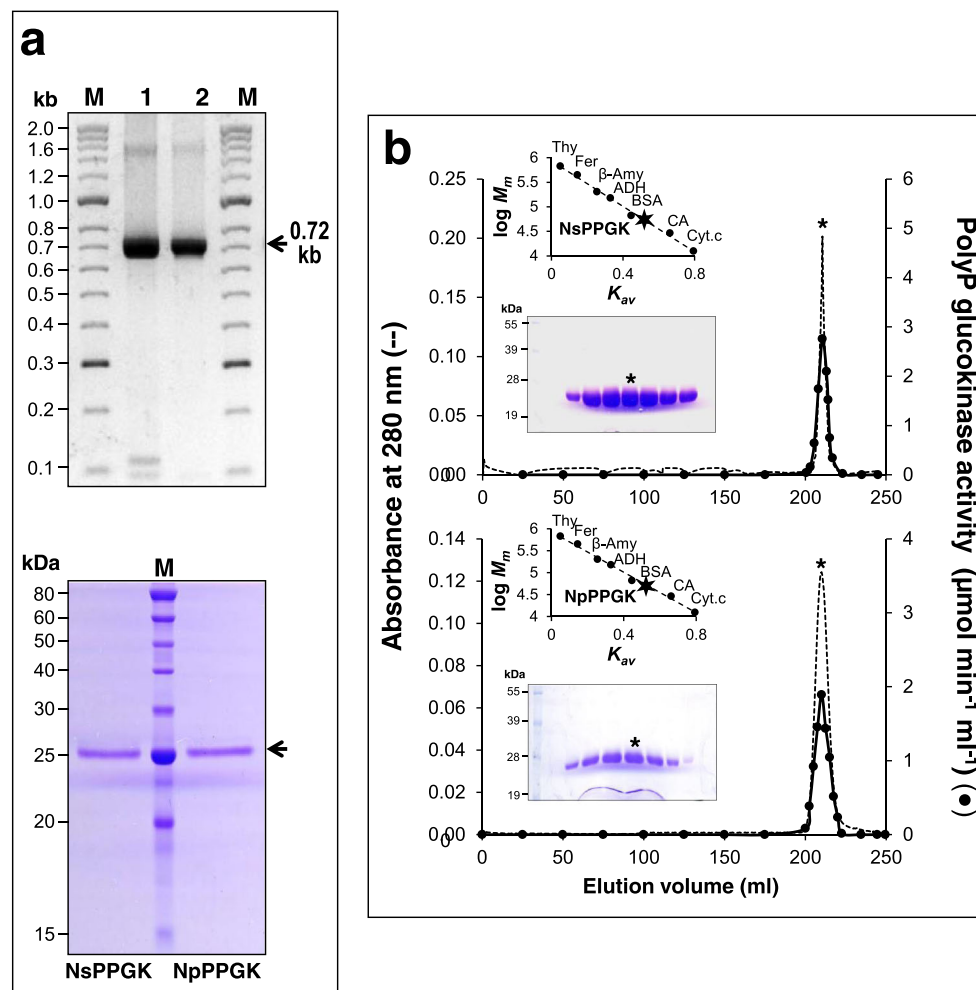


Fig. 1 **a** Upper panel. PCR amplification of cyanobacterial *ppgK* genes. An electrophoretic analysis of PCR-amplified DNA fragments corresponding to the *ppgK* genes of *Nostoc* sp. PCC7120 (lane 1) and *Nostoc punctiforme* PCC73102 (lane 2), and DNA size markers (M), is shown. Amplification reactions were performed with specific primer pairs and cyanobacterial genomic DNA as a template, as described in the “Materials and methods” section, and subsequently loaded onto 1.2 % agarose–TBE gel. As shown, a single DNA band of approximately 0.72 kb was obtained in each case (arrow). Lower panel SDS-PAGE (12 %, w/v, polyacrylamide; 0.5 %, w/v, SDS) analysis of recombinant NsPPGK and NpPPGK purified after FPLC gel filtration. Approx. 10 μg of NsPPGK (27.34 kDa, nominal subunit M_m) and NpPPGK (27.24 kDa, nominal subunit M_m) were applied per lane. M protein standards. Numerals on the left indicate the molecular masses (kDa). Arrow indicates the PPGK protein bands. **b** FPLC gel-filtration chromatography analyses of native M_m and oligomeric states of the PPGKs from *Nostoc* spp.

NpPPGK showed fairly high glucokinase activity levels with P_{13-18} as a substrate, in the range of 0.15 to 0.20 $\mu\text{mol min}^{-1} \text{mg}^{-1}$ protein, respectively. In contrast, no PPGK activity was detected in extracts from cells containing the pQE-80L plasmid with no insert. Milligram quantities of the respective N-terminal His₆-tagged fusion proteins were subsequently isolated in ca. 95 % purity after one-step affinity purification onto a HisTrap FF Crude Ni-NTA column (Online Resource Figs. S2 and S3, Table S4). Enzyme purity was

Aliquots (0.5 ml) of metal-chelated chromatography purified preparations of recombinant NsPPGK and NpPPGK were applied to a Superdex© 200 PG column. Calibration curves with protein standards (Thy thyroglobulin, Fer ferritin, Amy β -amylase, ADH alcohol dehydrogenase, BSA bovine serum albumin, CA carbonic anhydrase, Cyt.c cytochrome c) are displayed on the left upper corner of the chromatography elution profile figures. A SDS-PAGE analysis of selected fractions around the central peak fraction (50 μl aliquots applied per lane) is also shown. Note that single elution peaks, corresponding to absorbance at 280 nm (broken line) and polyP glucokinase activity (filled circles, solid line), overlapped in both cases. The asterisks indicate the fraction peaks of recombinant PPGKs as determined by their enzymatic activity and absorbance at 280 nm. Native M_m values of 49.4 and 55.1 kDa were estimated for NsPPGK and NpPPGK, respectively. K_{av} phase distribution coefficient of the analyzed proteins

further enhanced by following FPLC gel filtration chromatography, which was confirmed by electrophoresis on SDS-PAGE gels (Fig. 1, Online Resource Table S4). Thus, a single protein band of ca. 27 kDa was found in both purified PPGK preparations (Fig. 1a), in good agreement with the nominal M_m values of 27,339 and 27,236 Da predicted for the recombinant NsPPGK and NpPPGK polypeptides, respectively. Besides, native M_m values and oligomeric states of the recombinant proteins were determined by gel filtration chromatography,

and values of 49.4 ± 4 kDa and 55.1 ± 5 kDa (means $\pm$ SE of three independent determinations) were obtained for NsPPGK and NpPPGK, respectively (Fig. 1b). Therefore, both proteins adopted a stable dimeric arrangement in solution. In accordance with these results, MALDI-TOF determination of absolute M_n values gave values of $27,287 \text{ Da} \pm 0.1\%$ and $27,236 \text{ Da} \pm 0.1\%$ for the recombinant NsPPGK and NpPPGK subunits, respectively. In addition, the identities of the recombinant NsPPGK and NpPPGK polypeptides were confirmed by peptide mass fingerprinting covering respectively about 55 and 82 % of the natural sequences, and eventual identification by MALDI-TOF MS (Online Resource Fig. S4). Together, these active and high-purity fractions were used for the subsequent determination of their enzymatic kinetic parameters.

NsPPGK and NpPPGK are strictly polyP-dependent glucokinases with preference for long-chain PolyP

The purified recombinant NsPPGK showed no activity towards ATP, CTP, GTP, TTP, or dATP as compared to sorts of polyP (Fig. 2). The absolute specificity of NsPPGK for inorganic polyphosphates appears to be a common property of PPGK enzymes in other heterocystous filamentous cyanobacteria, since similar results were observed in the characterization of NpPPGK. The substrate specificities

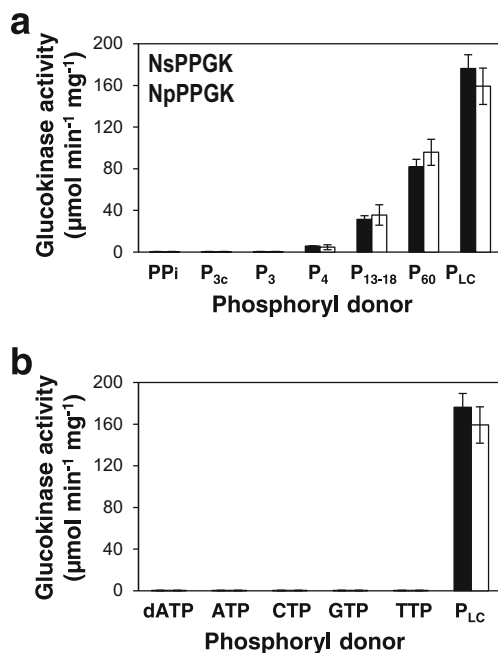


Fig. 2 Substrate specificity of cyanobacterial recombinant PPGKs. PolyP glucokinase activity levels of purified NsPPGK (black bars) and NpPPGK (white bars) were determined using polyPs of different chain lengths (a) or diverse NTPs (b) as phosphoryl donor substrates. Activity levels were obtained from three independent experiments and are shown as means $\pm$ SE. Note that both cyanobacterial PPGKs are strictly polyP-dependent glucokinases, and long-chain polyPs are their optimal substrates. No significant activity was detected with either NTPs, PPi, P_{3c}, or P₃

concerning polyP as phosphoryl donor to produce glucose 6-phosphate by cyanobacterial PPGKs were probed using synthetic polyP molecules of various chain lengths at saturating glucose levels (Fig. 2a). The rate of sugar phosphorylation for the polyP chain lengths followed a similar trend in both recombinant enzymes, longer polyP result in higher specific glucokinase activity. This indicated that PPGKs from cyanobacteria bound and hydrolyzed long-chain polyP substrates most efficiently (Table 1). This highlights its reasonable consistency with previous findings on polyP/ATP glucokinases of other bacteria (Girbal et al. 1989; Hsieh et al. 1996b; Tanaka et al. 2003; Mukai et al. 2003; Lindner et al. 2010a). Noteworthy, both cyanobacterial PPGKs are also able to use short-chain polyP. With reference to the sole crystal structure of a bacterial polyP/ATP glucomannokinase published to date (Mukai et al. 2004), it has been proposed that there is a minimal length between two phosphoryl groups consisting on a putative pentapolyphosphate binding site. However, NsPPGK and NpPPGK exhibit modest but significant specific activity levels with P₄ ($5\text{--}7 \mu\text{mol min}^{-1} \text{mg}^{-1}$ protein) (see Table 1). Analogous experiments revealed that cyanobacterial PPGKs were unable to use shorter polyP than P₄, such as P₃, P_{3c}, or PPi. Estimation of the kinetic parameters of NsPPGK with different polyP and hexose substrates revealed that there is a remarkable increase of the catalytic constant k_{cat} (more than 30-fold) with increasing polyP chain length from P₄ up to P_{LC}, while K_m values remain fairly constant, which explained the higher catalytic efficiency of long-chain polyP (Tables 1 and 2, Figs. S5 and S6).

To determine the mechanism of polyP utilization by NsPPGK, P₁₅₀ at saturation concentration was used as a phosphoryl substrate, while the progress of the reaction was monitored by collecting sequential aliquots at increasing times. PolyP were isolated and electrophoresed on a preparative polyacrylamide gel, and eventually visualized with toluidine blue O staining. As shown in Fig. 3, this medium-sized polyP was utilized by the cyanobacterial PPGK by an essentially non-processive mechanism, as was evidenced by the non-noticeable broadening of the range of polyP sizes with the reaction time. A non-processive mechanism is also consistent with the observed formation of polyP of intermediate sizes from the longest polyP during the reaction progress (see Fig. 3).

A variety of compounds which are analogs to the phosphoryl donors were also tested to assess whether or not they could act as PPGK activity inhibitors (Table 3). P₃ and PPi were fairly strong inhibitors, with K_i values of 0.13 and 0.19 mM, respectively, while ATP only modestly inhibited the activity of NsPPGK. In addition, a control experiment with increasing concentrations of NaCl was conducted to determine the effect of the ionic strength on the PPGK activity. Results from Table 3 indicated that NsPPGK was not as severely inhibited by NaCl as by short polyP or ATP, since the observed concentrations required for substantial enzyme inhibition, most

Table 1 Kinetic parameters of purified recombinant polyP gluco(manno)kinase from *Nostoc* sp. PCC7120 with different polyPs and hexoses as substrates

Substrate (polyP _n)	$V_{\max}$ ($\mu\text{mol min}^{-1} \text{mg}^{-1}$)	K_m^a (μM)	k_{cat} (s^{-1})	Catalytic efficiency k_{cat}/K_m ($\text{mM}^{-1} \text{s}^{-1}$)
P ₄	5.7	29.9	4.9	164
P _{13–18}	31.4	37.7	27.0	717
P ₆₀	81.7	39.8	70.2	1764
P _{LC}	176.3	49.5	151.4	3059
Glucose (P _{LC})	239.3	67.7	196.0	2895
Mannose (P _{LC})	3.1	2360	1.6	0.7

^a K_m values are calculated as polyP

probably produced by ionic strength effect, were much higher (50–150 mM range). An inhibitory effect on PPGK activity was also obtained with KCl being even more marked than that of NaCl (Table 3), thus suggesting that electrostatic forces may be involved in the interaction between polyP and the enzyme.

NsPPGK and NpPPGK possess a modest but significant polyP-dependent mannokinase activity

Besides glucose, the cyanobacterial PPGKs phosphorylated mannose as well, but just in a minor extent (Table 2). The specific activity values of NsPPGK for glucose and mannose were 229.1 and 3.1 $\mu\text{mol min}^{-1} \text{mg}^{-1}$, respectively. Values of the same order of magnitude were obtained for NpPPGK. A further study of their catalytic efficiencies evidenced that cyanobacterial PPGKs clearly exhibit a remarkable preference (approx. 100-fold higher) for glucose instead of mannose as a substrate (Table 2). Conversely, fructose was totally inactive as a phosphoryl acceptor.

Cyanobacterial PPGKs are divalent cation-dependent enzymes with distinctive alkaline pH optimum and remarkable thermotolerance

The activities of both cyanobacterial PPGKs were absolutely dependent on the presence of a divalent cation in the reaction

Table 2 PolyP-hexokinase activities of purified recombinant NsPPGK and NpPPGK

Hexose	Specific activity ($\mu\text{mol min}^{-1} \text{mg}^{-1}$)	
	NsPPGK	NpPPGK
Glucose (5 mM)	229.1 ± 14.0	174.3 ± 14.7
Mannose (50 mM)	3.1 ± 0.2	2.0 ± 0.1
Fructose (50 mM)	ND ^a	ND

ND not detected

^a The minimum level of detection was ca. 0.05 $\text{nmol min}^{-1} \text{mg}^{-1}$

mixture. Mg^{2+} was the optimal metal cofactor for glucose phosphorylation by both recombinant PPGKs, while Mn^{2+} and Fe^{2+} ions function in a lesser extent, and no activity was detected in the presence of Co^{2+} , Ca^{2+} , or Cu^{2+} ions (Fig. 4a). No activity was detected after incubation of the enzyme samples with 10 mM EDTA, and subsequent dialysis to remove all traces of EDTA resulted in complete loss of activity. Addition of 5 mM Mg^{2+} restored the full PPGK activity. The highest specific activity with magnesium ions was found in the concentration range 4–6 mM with an optimum estimated at 5 mM and a calculated K_m value of 1.3 mM (Fig. 4b). Higher concentrations of Mg^{2+} resulted in a decrease of PPGK activity.

Cyanobacterial PPGK activity was optimal at the alkaline pH range, between pH 8.5 and 9.0 (Fig. 5a). Nevertheless, the activity declined quite rapidly at higher pH values with no activity remaining at pH 10.5 or higher. A very similar pH dependence curve was obtained for both enzymes. Alkaline pH optimum is a common distinctive feature of other cyanobacterial enzymes when compared with their orthologs of non-photosynthetic bacteria and eukaryotes (Serrano et al. 1984; Serrano 1992).

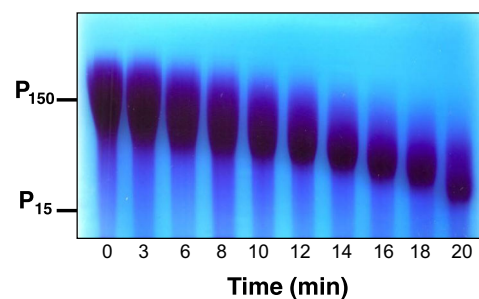


Fig. 3 Non-processive utilization of P₁₅₀ by NsPPGK; 2.5 mM of P₁₅₀ was used as a substrate for purified NsPPGK (approx. 3 $\mu\text{g/ml}$) following the standard assay conditions, as described in the “Materials and methods” section. At different time intervals, sequential aliquots were collected and polyP was isolated, electrophoresed on a preparative PAGE gel, and finally stained with toluidine blue. **Lane 1** is zero time, and **lanes 2 to 10** correspond to 3, 6, 8, 10, 12, 14, 16, 18, and 20 min, respectively

Table 3 Effect of different compounds on the activity of NsPPGK towards glucose and P_{13–18} as substrates

Chemicals	mM	Activity (%)
NaCl	1	100
NaCl	5	99.7
NaCl	10	99.5
NaCl	50	84.5
NaCl	150	66.0
KCl	1	95.8
KCl	5	82.3
KCl	10	70.4
KCl	50	65.2
KCl	150	49.9
ATP	1	96.3
ATP	5	71.0
ATP	50	12.2
PPi	1	99.8
PPi	5	55.6
PPi	10	28.4
P ₃	1	87.5
P ₃	5	39.9
P ₃	10	12.0
None		100

Specific activity in the absence of inhibitor (set as 100 %) was 31.8 $\mu\text{mol min}^{-1} \text{mg}^{-1}$

Both cyanobacterial PPGKs showed an optimal temperature as high as 45 °C (Fig. 5b). Indeed, PPGK from the actinobacterium *Arthrobacter* sp. (Mukai et al. 2003) exhibits a similar value, but considerably higher as compared to 30 °C for the PPGK from *M. phosphovor* and most other actinobacterial polyP/ATP glucokinases (Tanaka et al. 2003). To determine the thermostability of cyanobacterial PPGK, NsPPGK was preheated at 40, 50, 60, and 70 °C for 30 min. No loss of activity was observed after incubation below 50 °C. At 50 °C, 45 % of activity remained. However, only 7 % of PPGK activity remained at 60 °C, implying that NsPPGK is unable to tolerate these fairly high temperatures. Finally, this PPGK was irreversibly inactivated when exposed to temperatures above 60 °C for 30 min.

Closely related PPGK orthologs occur among diazotrophic *Cyanobacteria*

A bioinformatic search was carried out looking for PPGK homologs in a range of filamentous and unicellular strains representative of any of the five cyanobacterial taxonomic sections defined by Rippka et al. (1979). Then, a number of putative *ppgK* genes were identified in strains belonging to all these taxonomic groups. With two exceptions, the nitrogen-fixing strains *Synechococcus* sp. PCC7335 (Bergman et al. 1997) and *Synechococcus* sp. PCC7502, no PPGK-like ORFs were found so far among sequenced genomes of unicellular

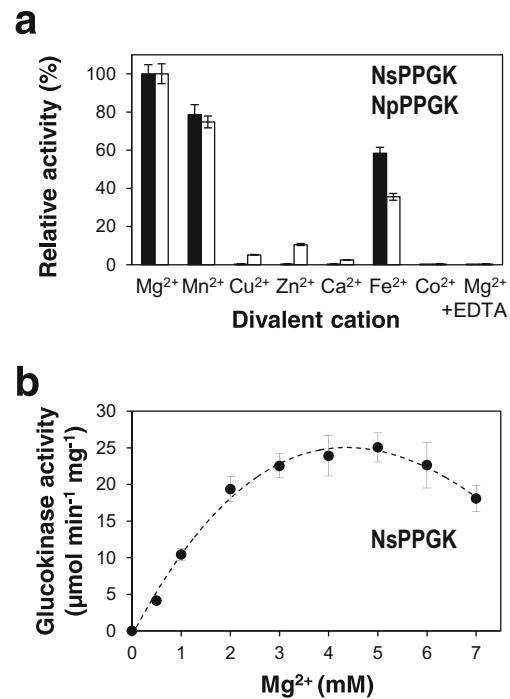


Fig. 4 Biochemical characterization of recombinant NsPPGK (black bars) and NpPPGK (white bars) regarding to metal cation dependence of polyP (P₆₀) glucokinase activity. **a** Metal cofactor specificity. Several divalent metal cations were added at 5 mM concentration to the assay mixtures. No detectable activity was measured with Mg²⁺ in the presence of 10 mM EDTA. Bars represent activity levels from three independent experiments and are shown as means±SE. Activity is expressed in relative units (100 % percentage assigned to the optimum condition in each case); 100 % activity levels correspond to 81.7±7.4 and 95.8±12.5 $\mu\text{mol min}^{-1} \text{mg}^{-1}$ for NsPPGK and NpPPGK, respectively. **b** NsPPGK activity dependence on Mg²⁺ concentration. Each point represents the mean activity value±SE of three independent experiments. As shown, no activity was detected either in the absence of a divalent cation or with an excess of the chelating agent EDTA

species from section I, typically non-nitrogen-fixing, e.g., *Synechocystis* sp. PCC6803 and *Synechococcus elongatus* PCC7942. Likewise, no hybridization band was observed in Southern blot analysis (Fig. 6) and no PPGK activity was detected in whole-cell extracts of the two latter strains (Table 4). In contrast, clear hybridization bands and significant levels of PPGK activity were detected with several polyPs in a number of strains of sections II to V with sequenced genomes exhibiting predicted *ppgK* genes as expected (such as *Dermocarpa* sp. PCC7437 and *Nostoc* spp.), but also in other diverse *Cyanobacteria* whose genomes have not been sequenced yet, such as the section III strain *Pseudanabaena* sp. PCC6903; the section IV strains *Nostoc* sp. PCC6719, *Calothrix* sp. PCC7601, *Calothrix* sp. PCC9327, *Anabaena* sp. ATCC33047, and *Nodularia chucula*; and section V strain *Fischerella muscicola* (Fig. 6, Table 4). Therefore, closely related putative polyP-dependent PPGKs seem to be widely distributed among diazotrophic, mostly multicellular, cyanobacterial strains.

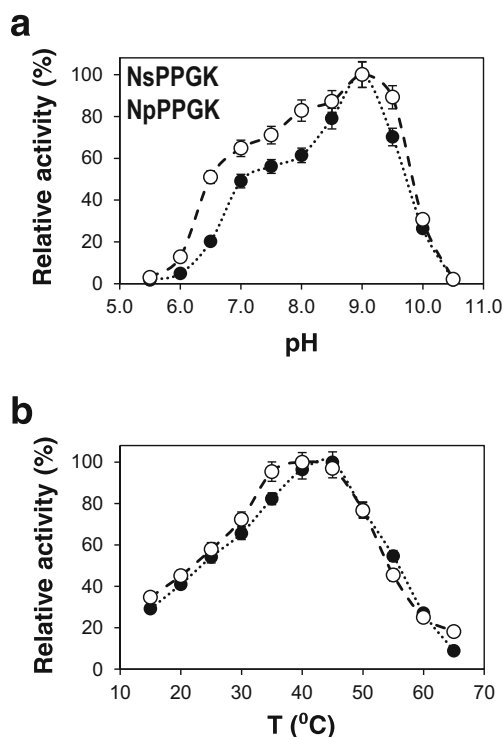


Fig. 5 Effect of the pH (**a**) and temperature (**b**) on the polyP (P_{60}) glucokinase activity of NsPPGK (filled circles) and NpPPGK (open circles). Data are shown as relative units (100 % percentage) and were assigned to the optimum condition in each case. Activity levels were obtained from three independent experiments and are shown as means $\pm$ SE; 100 % activity values correspond to 80.6 ± 7.3 and $93.5 \pm 9.2 \mu\text{mol min}^{-1} \text{mg}^{-1}$ (**a**) and 80.4 ± 6.7 and $92.7 \pm 7.3 \mu\text{mol min}^{-1} \text{mg}^{-1}$ (**b**) for NsPPGK and NpPPGK, respectively

Discussion

A BLAST sequence similarity search in the genome of *Nostoc* sp. PCC7120 revealed one ORF, *all1371*, with high homology to the well-characterized *M. tuberculosis* H37Ra *ppgK* gene (Szymona and Widomski 1974; Hsieh et al. 1996a). A similar approach revealed another putative PPGK-encoding gene, *Npun_R1878*, in the genome of *N. punctiforme* PCC73102, which was annotated to be encoding a ROK (transcriptional regulator/sugar kinase) family protein which shares a 91 % sequence identity with its homolog of *Nostoc* sp. PCC7120. Subsequent searches in bioinformatic databases identified about other 40 putative cyanobacterial PPGK orthologs. They are predicted to be highly similar proteins of about 230–250 amino acid residues, clearly smaller than their conventional ATP glucokinase counterparts (290–330 residues), and most of them were unprecisely annotated as ROK family proteins or transcriptional regulators/sugar kinases. This finding together with the deduced ROK family domain architecture characteristic of other previously reported bacterial PPGKs predicted for all cyanobacterial orthologs, led us to investigate whether glucose 6-phosphate synthesis could take place in *Cyanobacteria* through PPGK enzymes.

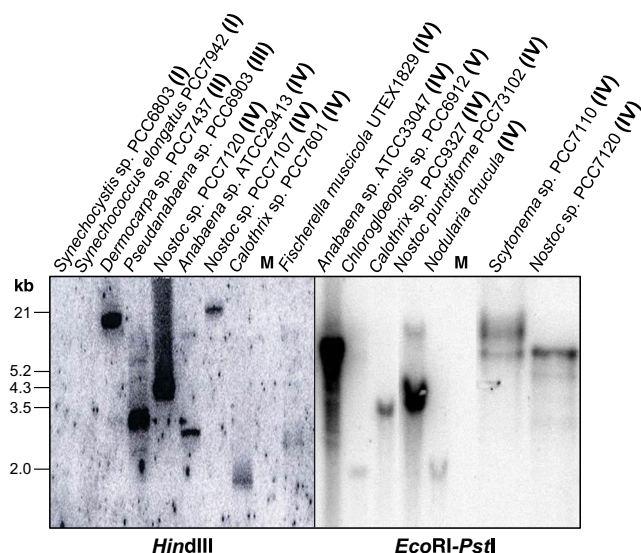


Fig. 6 Experimental evidence for the widespread occurrence of homologs of *Nostoc* spp. *ppgK* genes among diazotrophic *Cyanobacteria*. A search of putative *ppgK* genes was carried out by Southern blot analysis with diverse cyanobacterial strain representatives of the taxonomic sections (Roman numerals) in the classification of Rippka et al. (1979). The strains are identified by their collection numbers. The positions of *EcoRI*–*HindIII*-restricted λ DNA fragments used as standards (in the range of 21 to 2 kb) are indicated on the left side. Genomic DNAs (approx. 5 μg) were digested with *HindIII* (left panel) or *EcoRI* (right panel) restriction enzymes. The full coding sequence of the *ppgK* gene from *Nostoc* sp. PCC7120 was used as a probe under heterologous hybridization conditions at 55 °C. As shown, no hybridization band was observed only in the lanes corresponding to unicellular non-diazotrophic *Cyanobacteria* *Synechocystis* sp. PCC6803 and *Synechococcus* sp. PCC7942

This presumption was confirmed by the biochemical characterization of two recombinant cyanobacterial PPGK proteins purified by metal-affinity and size-exclusion chromatographies as described above. As a result, both *Nostoc* proteins were functionally validated with the ability to phosphorylate glucose and, to a lesser extent, mannose. These enzymatic reactions occurred using a wide range of polyP with different chain lengths as phosphoryl donors. However, no activity was detected with the shortest-chain-length polyPs, namely pyrophosphate (PPi), P_{3e} , or P_3 . In fact, P_4 was confirmed as the shortest polyP active as a substrate for PPGK enzymes described so far. Concerning the length of the chain of active polyP substrates, both *Nostoc* proteins seemed to follow a similar pattern, with higher catalytic efficiencies for long-chain polyP ($\geq \text{P}_{60}$). Remarkably, reaction rates of polyP utilization increased considerably with the number of phosphate residues per molecule. To our knowledge, Mukai et al. reported the sole crystal structure so far available for a polyP/ATP glucokinase—from the actinobacterium *Arthrobacter* sp. KM—and it was complexed with one glucose and two phosphate molecules instead of polyP (Mukai et al. 2004). According to this model, P_5 has been claimed as the shortest polyP able to enzymatically phosphorylate glucose. In contrast, we showed that both NsPPGK

Table 4 PPGK-specific activity levels with different polyP substrates in whole-cell extracts from diverse *Cyanobacteria*

Cyanobacterial strain	P ₄	P _{13–18} (nmol ^{−1} min ^{−1} mg ^{−1})	P ₆₀	P _{LC}
<i>Synechococcus elongatus</i> PCC7942(I)	ND ^a	ND	ND	ND
<i>Synechocystis</i> sp. PCC6803 (I)	ND	ND	ND	ND
<i>Dermocarpa</i> sp. PCC7437 (II)	9.0±0.7	8.4±0.7	8.5±0.7	10.9±1.3
<i>Pseudanabaena</i> sp. PCC6903 (III)	3.6±0.4	15.1±1.3	11.1±0.9	16.3±1.4
<i>Anabaena</i> sp. ATCC29413 (IV)	6.8±0.7	27.4±1.7	18.2±1.5	16.0±1.8
<i>Calothrix</i> sp. PCC9327 (IV)	0.1±0.3	2.1±0.2	3.1±0.2	4.6±0.3
<i>Nostoc punctiforme</i> PCC73102 (IV)	10.5±0.9	15.4±1.2	13.7±1.1	19.0±1.4
<i>Nostoc</i> sp. PCC7120 (IV)	0.8±0.1	17.9±1.3	18.4±1.5	10.9±0.9
<i>Scytonema</i> sp. PCC7110 (IV)	0.4±0.1	5.7±0.8	3.1±0.5	12.8±1.6
<i>Chlorogloeopsis</i> sp. PCC6912 (V) ^b	ND	ND	ND	ND
<i>Fischerella muscicola</i> UTEX1829 (V) ^b	ND	ND	ND	ND

Roman numerals in parentheses indicate the sections of the taxonomic classification of Rippka et al. (1979). The two unicellular strains of section I were grown in the presence of 2 mM NaNO₃ as a nitrogen source

ATCC American Type Culture Collection, PCC Pasteur Culture Collection, UTEX University of Texas at Austin Culture Collection, ND not detected activity

^a The minimum level of detection was ca. 0.05 nmol min^{−1} mg^{−1}

^b The presence of large amounts of extracellular mucous material made the achieving of reliable measurements with these colonial strains very difficult

and NpPPK are able to generate glucose 6-phosphate when using P₄ as a phosphoryl donor, although with a lower efficiency than longer-chain polyPs. Apart from this, it is noteworthy that both cyanobacterial PPGKs were strictly dependent on polyP, as there was no activity detected when ATP or any other NTP were used as phosphoryl substrates. This feature has been only described to date for the PPGK of the primitive, polyP-accumulating actinobacterium *M. phosphovor* (Tanaka et al. 2003). Here, we describe a novel subfamily of PPGK enzymes characteristic of *Cyanobacteria*, all of them being strictly dependent of polyP as the phosphoryl donor.

Using multiple sequence alignment of the polyP/ATP glucomannokinase from *Arthrobacter* sp. KM and other actinobacterial polyP/ATP glucokinases, a specific extra heptapeptide (PEAPAAAG) was identified in the conserved glucose region of the former protein which was proposed as responsible for the mannose-phosphorylating ability of the polyP/ATP glucomannokinase. In fact, PPGK from *Arthrobacter* sp. KM can phosphorylate fructose as well (Mukai et al. 2003). In addition, Szymona et al. have shown that when *M. phlei* was grown on fructose, a polyP-fructokinase activity was found. Contrastingly, when grown on mannose, polyP mannokinase was detected (Szymona and Ostrowski 1964). Nonetheless, despite lacking such heptapeptide, this work shows that both cyanobacterial PPGKs are able to phosphorylate mannose, although with fairly modest levels and a notably reduced catalytic efficiency compared to glucose. However, no significant polyP-fructokinase activity was detected for NsPPGK and NpPPGK.

Some other features of NsPPGK and NpPPGK were in some extent distinct to those previously described for other bacterial PPGKs. Thus, optimum pH was clearly alkaline, 8.5–9.0, while actinobacterial PPGKs have almost neutral optimal pH values (e.g., 7.5 for the *Arthrobacter* enzyme). Also, the notable thermostability of cyanobacterial PPGKs (optimal temperature, ca. 45 °C) is an outstanding catalytic feature that, like its alkaline optimal pH, may have biotechnological relevance. Other biochemical features were similar to those of other sugar kinases; thus, both cyanobacterial PPGKs required divalent metal cations, to which Mg²⁺ was preferred. Similarly, they were identified as homodimers although with natural subunit *M_m* values somewhat lower than those of bacterial polyP/ATP-dependent glucokinases (ca. 30 kDa) and eukaryotic hexokinases (ca. 35 kDa).

It has been hypothesized that polyP could be the phosphoryl donors for ancient organisms, and they were later replaced by ATP in the evolution (Lipmann 1965). This is based on the assumption that the Gibbs free energy of polyP hydrolysis is similar to the ATP, and their likely occurrence since prebiotic times. An interesting observation is that PPGK activities have been reported to date only in the comparatively ancient order of *Actinomycetales*. Noteworthy, in bacteria belonging to this order, the ratio polyP glucokinase vs. ATP glucokinase activities is higher in more phylogenetically ancient representatives (Hsieh et al. 1993; Phillips et al. 1999). According to this hypothesis, the following stage in the evolution of sugar kinases might be played by the dual ATP/polyP glucokinases, like the PPGKs described in *Propionibacterium shermanii*, *M. tuberculosis*, or other *Actinobacteria* (Pepin and Wood

phylogenetic data shown, their encoded PPGK proteins form a compact well-supported cluster, clearly divergent from the actinobacterial PPGK assembly, with a number of putative orthologs encoded by the genomes of selected unicellular, colonial, and filamentous cyanobacterial strains. It should be noted in this respect that about 50 putative cyanobacterial PPGK orthologs were identified in database searches (July 2014) (Online Resource Table S3). Noteworthy, PPGK orthologs of marine cyanobacterial strains such as *Acaryochloris marina* MBIC11017 and *Nodularia spumigena* CCY9414 are also included in this group. This suggests that cyanobacterial polyP is possibly used as an alternative source of energy in place of ATP in the ocean environment as well. All in all, our sequence comparison and molecular phylogenetic data reveal that cyanobacterial PPGKs are structurally simpler and presumably more ancient than their homologs of *Actinobacteria*. With the exception of the enzyme of the polyP-accumulating actinobacterium *M. phosphovorus*, all the other PPGKs described to date utilize polyP as well as ATP. These findings agree with the ancestral character of *Cyanobacteria* and suggest that the strictly polyP-dependent PPGKs may represent molecular relicts of a hypothetical ancient world in which polyP could be preferentially used for metabolic functions.

The present work also envisages new perspectives for an innovative cost-effective enzymatic production of glucose 6-phosphate or mannose 6-phosphate by a novel class of strictly polyP-dependent glucokinases from diazotrophic *Cyanobacteria*. Thus, synthesis of sugar phosphates could be performed by immobilized-engineered cyanobacterial PPGKs from the very stable inorganic polymer polyP without continuous regeneration of ATP, an expensive cofactor required by conventional hexokinases.

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