

Autoregulation of *fixK₂* gene expression in *Bradyrhizobium japonicum*

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Abstract Several essential *Bradyrhizobium japonicum* genes for a symbiotic, nitrogen-fixing root-nodule symbiosis are positively controlled under micro-oxic conditions by the FixLJ–FixK₂ regulatory cascade. Negative control is exerted by reactive oxygen species at the level of the FixK₂ protein. Furthermore, we noticed that *fixK₂* gene expression is increased in a *fixK₂* mutant, suggesting that FixK₂ in the wild type has a negative effect, directly or indirectly, on its own expression. To possibly understand this effect, the transcription pattern of the *fixLJ-bll2758-fixK₂* gene region was examined more closely. While *fixK₂* gene transcription is activated by FixJ, the *bll2758* gene is transcribed from its own promoter in a FixK₂-dependent manner, and there is no read-through transcription from *bll2758* into *fixK₂*. The *bll2758*-encoded protein is predicted to be a stand-alone receiver domain of a response regulator, making it a prime candidate for exerting an inhibitory role on the expression of *fixK₂*. Transcriptome profiling of a *bll2758* knock-out mutant revealed, however, that neither *fixK₂* itself nor any of the known FixJ- and FixK₂-dependent target genes is significantly affected in their expression. This precludes a role of the *bll2758* product as a so-called FixT-like protein in the inhibition of FixLJ function, as was proposed for *Sinorhizobium meliloti* and *Caulobacter crescentus*. Instead, we rationalize that other transcription factors, whose genes are activated by FixK₂, might be involved in the negative autoregulation of *fixK₂* gene expression.

Keywords *Bradyrhizobium japonicum* · Gene regulation · Microarrays · Promoter · Transcription

Abbreviations

cDNA Complementary DNA
FixJ-P Phosphorylated FixJ
ORF Open reading frame
PCR Polymerase chain reaction

Introduction

Strong induction of bacterial gene expression is often achieved either when target genes are subject to dual positive control, involving two synergistically acting, disparate activators, or when two or more transcription activators work sequentially in a cascade in which the first induces expression of the gene for the second, thus boosting the level of the ultimate activator of target genes. These arrangements also allow the integration of additional regulatory cues at different levels. The FixLJ–FixK cascade, addressed in this work, which exists in many symbiotic, nitrogen-fixing rhizobia, is an example of a sequentially acting pathway (for reviews, see Fischer 1994; Batut and Boistard 1994; Dixon and Kahn 2004). FixL, a sensor-histidine autokinase, responds to a decreased oxygen tension and subsequently phosphorylates the response regulator FixJ. FixJ-phosphate (FixJ-P) then activates the gene for FixK, a member of the CRP/FNR protein family. FixK, in turn, activates genes and operons related to micro-oxic energy metabolism, e.g., the *fixNOQP* and *fixGHIS* operons for the synthesis and function of the high-affinity *cbb₃*-type oxidase.

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In contrast to the gene activation steps in the FixLJ-FixK cascade, less information is available about how the genes are down-regulated. Four negatively interfering modes of control have been described for either *Sinorhizobium meliloti* or *Bradyrhizobium japonicum*: (1) decrease of the FixJ-P level by oxygenation of FixL (Lois et al. 1993; Sousa et al. 2007); (2) negative autoregulation of *fixK* gene expression (Foussard et al. 1997; Nellen-Anthamatten et al. 1998; Garnerone et al. 1999); (3) inhibition of *fixK* gene expression by a precursor of purine biosynthesis (Soberón et al. 2001) and; (4) posttranslational control of FixK₂ activity by reactive oxygen species (Mesa et al. 2009). The mechanism of *fixK* autoregulation in *S. meliloti* is indirect (Fig. 1a). The FixK protein activates the *fixT*₁ gene, encoding a stand-alone receiver-domain protein that was shown to inhibit FixL autophosphorylation (Foussard et al. 1997; Garnerone et al. 1999). The anti-kinase activity of FixT₁ seems to be assisted by an asparagine synthetase-like protein (AsnO), but the physiological meaning of this kind of control remains to be elucidated (Bergès et al. 2001). Many elements of the *S. meliloti* *fixK* autoregulation mechanism were found to operate in a similar way in the non-symbiotic, nitrogen non-fixing *Caulobacter crescentus* (Crosson et al. 2005; Paul et al. 2008; Jenal and Galperin 2009), although the precise site of inhibition by FixT is not known (Fig. 1b). A *fixT*-like gene (*bll2758*) was also identified previously in *B. japonicum* (Nellen-Anthamatten et al. 1998; Kaneko et al. 2002), where it is located between the *fixLJ* and *fixK*₂ genes (Fig. 1c). The aim of this work was to unravel the transcriptional organization of the *bll2758*-*fixK*₂ region and to find out whether the *bll2758* product functions like the FixT protein. The result was that *B. japonicum* differs in this respect from *S. meliloti* and *C. crescentus*. Although we could demonstrate FixK₂-dependent transcription activation of the *bll2758* gene, the product of the latter does not appear to interfere with the expression of either *fixK*₂ or known FixK₂-dependent genes. An alternative scenario for the negative autoregulation of *fixK*₂ is discussed.

Materials and methods

Media and growth conditions

Bradyrhizobium japonicum strains were grown either oxidically with vigorous shaking or micro-oxidically (0.5% O₂ and 99.5% N₂ in the gas phase) in a modified peptone-salts-yeast extract medium (Mesa et al. 2008). The growth temperature in all experiments was 30°C. Where appropriate, cultures contained the following concentrations of antibiotics (in µg per mL): spectinomycin, 100; kanamycin, 100. *Escherichia coli* strains harboring recombinant plasmids

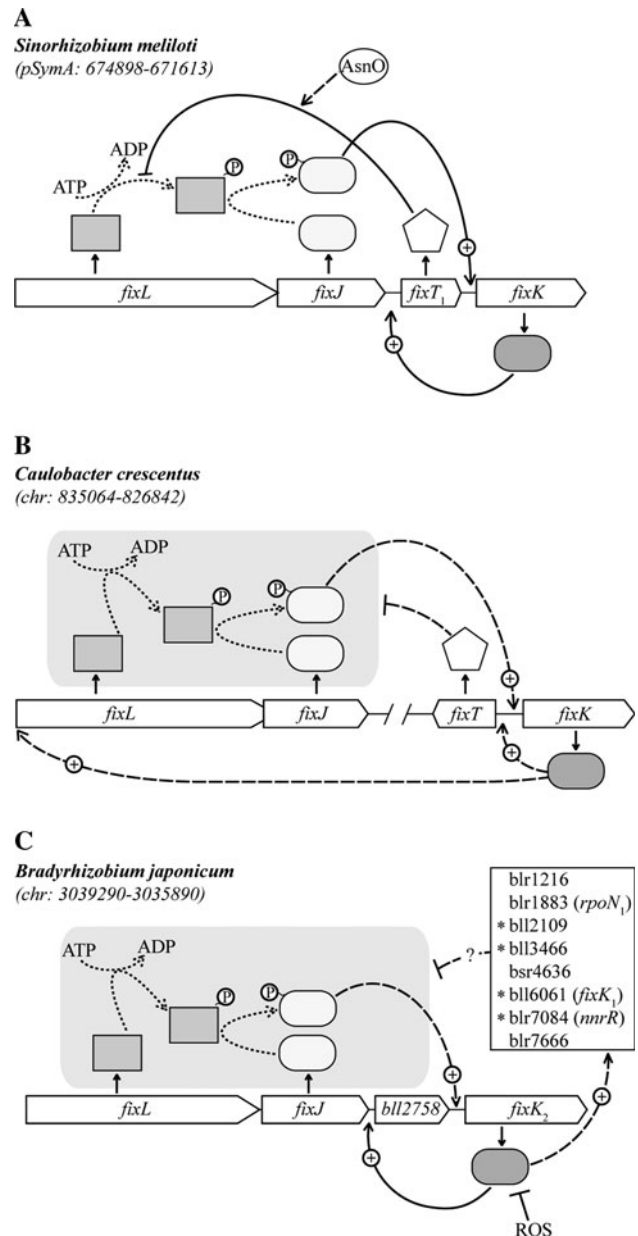


Fig. 1 Comparison of FixLJ-FixK regulatory circuits in *S. meliloti* (a), *C. crescentus* (b), and *B. japonicum* (c). The scheme for *B. japonicum* reflects work elaborated not only previously but also in this work. See text for a description of the three models. Genes are drawn to scale. Their genome coordinates are shown below species names. Specific geometric shapes are used for homologous gene products. The large gray-shaded rectangle in b and c means that, in contrast to a, the precise site of negative autoregulation is not known. The stand-alone receiver domain encoded by *fixT*/*bll2758* appears to play a negative regulatory role only in a and b. Instead, other transcription factors compiled in a box may play this role in c. Of these, CRP/FNR family members are marked with an asterisk. Solid lines: biochemically proven direct interactions. Dashed lines: direct or indirect interactions. Dotted lines: phosphoryl group transfer. Lines ending with a circled + and/or arrowheads symbolize positive control. Lines ending with a T bar symbolize negative control. ROS, reactive oxygen species

were grown in Luria-Bertani medium at 37°C (Miller 1972). Antibiotics were used as follows (in µg per mL): ampicillin, 200; kanamycin, 30.

Bradyrhizobium japonicum strains

Bradyrhizobium japonicum strains used in this work were 110*spc4* (wild type, Regensburger and Hennecke 1983); 7360 (*fixJ::aphII*, Anthamatten and Hennecke 1991); 9043 (Δ *fixK*₂, Nellen-Anthamatten et al. 1998); 7454 (*fixK*₁::*aphII*, Anthamatten et al. 1992); 7414 (*bll2758::aphII*, Anthamatten and Hennecke 1991); 8678 (Δ *nnrR*, Mesa et al. 2003); and N50-97 (*rpoN*₁::*aphII* + *rpoN*₂:: Ω , Kullik et al. 1991). Insertion of a translational *fixK*₂'-*lacZ* fusion into the chromosomes of the wild type, the Δ *fixK*₂ mutant, and the Δ *nnrR* mutant gave rise to strains 9054 (Nellen-Anthamatten et al. 1998), 9054K2 (Nellen-Anthamatten et al. 1998), and 9054R (this work), respectively. β -Galactosidase activity in these strains was determined as described previously (Fischer et al. 1993).

Reverse transcription PCR

The technique was used to determine if the two adjacent genes *bll2758* and *fixK*₂ are transcribed as an operon. RNA template from micro-oxically grown *B. japonicum* wild-type cells was reverse-transcribed to cDNA with the same method as used in microarray sample preparation (Hauser et al. 2007). Subsequently, the cDNA was used as template for a hot-start PCR with *Pfu* DNA polymerase. The conditions were 30 s at 94°C for melting; 60 s at 60°C for annealing; and 90 s at 68°C for elongation. 35 cycles were run. The following primers were employed in different pairwise combinations: LR15, 5'-TCTCCTCGTCCTTGGGTTAG; LR16, 5'-TAGCCGGTGATCAGGATCACG; LR17, 5'-CAACGGAGATGGCGCAGATG; LR19, 5'-GCTGGAAACCGACGGCTTTG. Their positions and orientations are depicted in Fig. 2.

In vitro transcription

Assays for multiple-round in vitro transcription were carried out in a volume of 20 µl under standard conditions as described previously (Beck et al. 1997; Mesa et al. 2005). Plasmids used as transcription templates were based on pRJ9519 which contains the *B. japonicum* *rrn* terminator (Beck et al. 1997). A 210-bp *EcoRI* fragment with the *bll2758* promoter region was cloned into pRJ9519, resulting in plasmid pRJ0208. The *fixN* promoter-containing plasmid pRJ8816 (Mesa et al. 2005) was applied as a positive-control template. Purified FixK₂ protein (Mesa et al. 2005) was used in amounts of 1.25 or 2.5 µM. RNA size markers were prepared as described (Mesa et al. 2005).

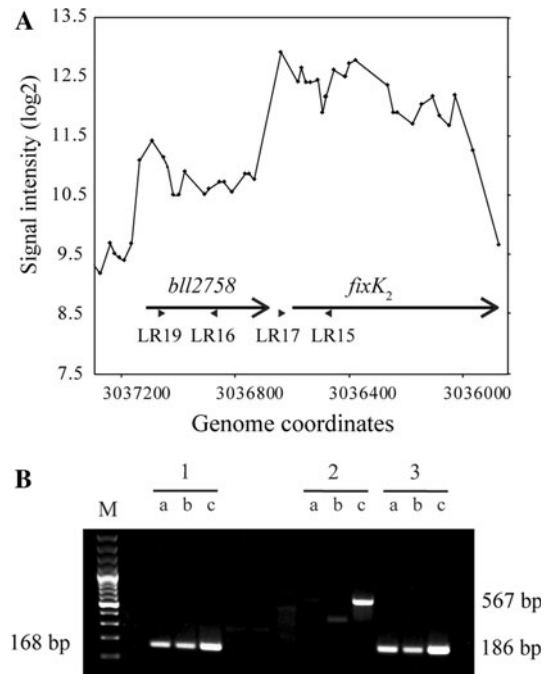


Fig. 2 Transcriptional organization of the *bll2758*–*fixK*₂ locus in *B. japonicum*. **a** Signal intensities derived from different oligonucleotide probes of the *bll2758*–*fixK*₂ region on the Gene Chip microarray. Data were obtained with RNA from micro-oxically grown wild-type cells. **b** Reverse transcription PCR. LR15, LR17, LR16, LR19 plus the associated small, filled arrowheads denote the binding sites and orientations of primers. cDNA was obtained from RNA of microoxically grown wild type. Lanes labeled with *a*, 50 ng cDNA; lanes labeled with *b*, 5 ng cDNA; lanes labeled with *c*, 50 ng genomic DNA (control). Primer combinations are indicated above the gel: 1 LR15 + LR17, 2 LR15 + LR19, 3 LR16 + LR19. Lengths (in bp) are indicated next to the respective band. *M* markers

A phosphoimager (Imager FX Pro; Bio-Rad Laboratories, Hercules, CA, USA) was used to detect the transcripts. For visualization, Quantity One Software, Version 4.6.7 (Bio-Rad Laboratories), was applied.

Primer-extension experiments

The transcription start site of *bll2758* was determined with methods reported previously (Beck et al. 1997; Mesa et al. 2005). Primer 9519-1 (5'-ATGGATGAACTCCAAGACGGTATC) was used for primer extension with the transcript synthesized in vitro from plasmid pRJ0208. To determine the transcription start site of *bll2758* in vivo, RNA template was extracted from micro-oxically grown *B. japonicum* wild-type cells, and primer LR21 (5'-GATCCCAGGACTGCGGCGTCATCATC) was used for extension.

Microarrays

Genome-wide transcription levels were determined as detailed previously, using a custom-designed *B. japonicum*

Affymetrix Gene Chip (Hauser et al. 2007; Mesa et al. 2008, 2009). In this work, the transcription profile of strain 7414 (*bll2758::aphII*) was analyzed. The complete dataset has been deposited in the Gene Expression Omnibus database (GEO; <http://www.ncbi.nlm.nih.gov/geo>) under GEO Series accession number GSE21400. Tiling analysis was done as described by Hauser et al. (2007) using the Affymetrix Tiling Analysis Software (Affymetrix, Santa Clara, CA, USA).

Bioinformatics

Potential FixK₂ binding sites in DNA sequences were identified by searching for TTG-N₈-CA motifs and by searching for sequences containing the TTG stretch and at least three additional nucleotides of the consensus TTGAT-N₄-ATCAA (Mesa et al. 2005). The program T-COFFEE (<http://www.ebi.ac.uk/Tools/t-coffee/index.html>) was used for the alignment of the predicted 118-amino-acid Bll2758 protein with other stand-alone receiver domains. The result was then visualized with GeneDoc 2.7 (<http://www.psc.edu/biomed/genedoc>).

Results

The *bll2758* and *fixK₂* genes are not cotranscribed

The organization of the *B. japonicum fixL-fixJ-bll2758-fixK₂* gene region is shown in Fig. 1c. The Kazusa nomenclature is used for *bll2758* (<http://genome.kazusa.or.jp/rhizobase>) because, for reasons given below, the previous designation ORF138 (Anthamatten and Hennecke 1991) was found to be inappropriate. A FixJ-P-activated promoter is located in front of *fixK₂* (Nellen-Anthamatten et al. 1998). It was noticed previously that expression of a chromosomally integrated *fixK₂'-lacZ* fusion led to a more than tenfold higher β -galactosidase activity in a *fixK₂* knock-out mutant as compared with the wild-type background, when cells of both strains had been grown under micro-oxic conditions (Nellen-Anthamatten et al. 1998). This observation was confirmed after inspection of data recently obtained by microarray-assisted transcriptome analyses: again, transcription from the *fixK₂* promoter was found to be up-regulated in a *fixK₂* background (Mesa et al. 2008). This suggested that FixK₂ regulates its own structural gene, directly or indirectly, in a negative way. Therefore, as a first approximation, the DNA upstream and around the beginning of *fixK₂* was examined for the presence of FixK₂ binding sites, using the canonical FixK₂ box (5'-TTGAT-N₄-ATCAA-3') as the query sequence (Mesa et al. 2005, 2008), although we did not know whether a repressor-binding site would look similar to the *bona fide*

activator binding site. One sequence (5'-TTGAG-N₄-GTCAA-3') that complied best with the consensus was identified at the beginning of *bll2758*. We reasoned that FixK₂ bound to that site could exert a negative effect on *fixK₂* expression only if there was a read-through transcription from *bll2758* into *fixK₂*. The results of experiments shown in Fig. 2 clearly demonstrate that this is not the case. The tiling-like architecture of the Gene Chip used (Hauser et al. 2007) helped reveal a non-uniform transcription intensity throughout this region, with a marked increase of transcription at the start of *fixK₂* (Fig. 2a). More importantly, using reverse transcription PCR, we did not detect RNA spanning the *bll2758* and *fixK₂* genes (Fig. 2b, lanes 2a and 2b, with the primer pair LR15/LR19), whereas individual transcripts from within *fixK₂* (lanes 1a and 1b) and *bll2758* (lanes 3a and 3b) were detected. Hence, *bll2758* and *fixK₂* are transcribed as separate genes, not as an operon.

The *bll2758* gene is a new direct target for activation by FixK₂

Although the putative FixK₂ box described above was very near the translational start of the annotated *bll2758* ORF, we suspected that it might rather be a promoter element than a repressor binding site. To clarify the issue, the possible activation of the *bll2758* gene by FixK₂ was first tested in a cell-free transcription assay (Mesa et al. 2005) with *B. japonicum* RNA polymerase holoenzyme and purified FixK₂, and using the *bll2758* 5'-region as DNA template. The result is shown in Fig. 3. A *bll2758*-specific transcript was synthesized only when the FixK₂ protein was present in the assay (lanes 2 and 3). Accordingly, this gene is a newly identified direct target for transcription activation by FixK₂, thus expanding the list of 11 approved FixK₂ targets (Mesa et al. 2008) by one additional member. Next, the transcription start site of *bll2758* was determined by primer extension (Fig. 4). Both, the in vitro synthesized RNA and the RNA extracted from micro-oxically grown cells gave rise to an extension product whose end mapped to a G (position +1; Fig. 4a, b) just two nucleotides before the annotated start codon of the *bll2758* ORF (Fig. 4c). This is too close to accommodate a ribosome binding site. Most likely, therefore, the true translation start is 20 codons further downstream at a GTG triplet (Fig. 4c), and the ORF contains 118 instead of 138 codons. ORF138, the name coined previously for *bll2758* (Anthamatten and Hennecke 1991; Nellen-Anthamatten et al. 1998), was henceforth abandoned. The -10 and -35 regions upstream of the transcription start site possess reasonably well conserved promoter sequences. Of particular interest is the partial overlap of the -35 region with the aforementioned FixK₂ box, as depicted in Fig. 4b. This arrangement is

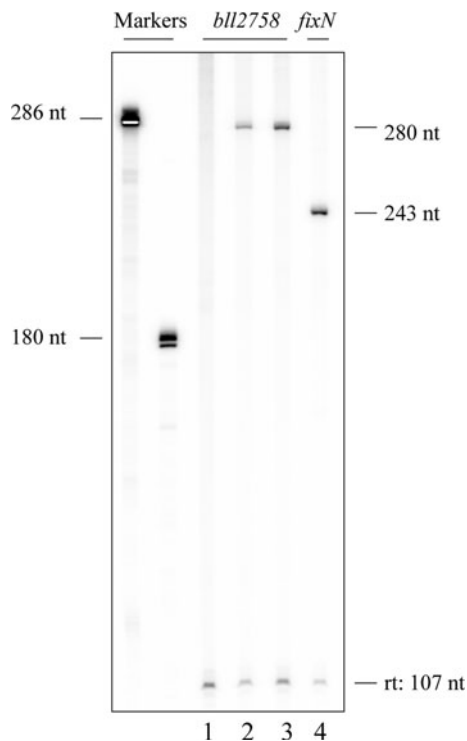


Fig. 3 In vitro transcription from the *bll2758* promoter. Transcripts from template plasmid pRJ0208, which contains the *bll2758* promoter region, were generated by multiple-round in vitro transcription with purified *B. japonicum* RNA polymerase holoenzyme and purified FixK₂ protein. Plasmid pRJ8816, which contains the promoter of the FixK₂-dependent *fixN* gene was used as control. Markers were generated as described previously (Mesa et al. 2005). FixK₂ protein concentrations were as follows: lane 1, no protein added; lanes 2 and 4, 1.25 μM; lane 3, 2.5 μM. The positions of the FixK₂-dependent transcripts, the FixK₂-independent vector-encoded reference transcript (rt), and the two RNA size markers are labeled; nt nucleotides

characteristic for all promoters of FixK₂-activated genes (Mesa et al. 2008). Taken together, FixK₂ positively controls *bll2758* (Fig. 1c), in a similar way as *fixT* is regulated by FixK in *S. meliloti* and *C. crescentus* (Fig. 1a, b).

Unlike FixT, the *bll2758* product is not a *fixK₂*-specific negative regulator

With the newly proposed translational start at the GTG codon (Fig. 4c), the *bll2758* gene codes for a predicted 118-amino acid protein that shares a low, but significant sequence similarity with previously described FixT-like single-domain response regulators (Jenal and Galperin 2009) of different proteobacteria (Fig. 5). Among several functionally conserved amino acids is a strictly conserved aspartic acid residue as the site of phosphorylation (Fig. 5). Being devoid of DNA-binding domains, such proteins do not directly affect transcription but modulate gene expression indirectly by interacting with other regulators. *S. meliloti* FixT₁, for example, inhibits FixL autokinase

activity (Foussard et al. 1997; Garnerone et al. 1999). To unravel any effect on the expression of *fixK₂* and FixK₂-dependent genes, the *bll2758* mutant (strain 7414) grown micro-oxically was investigated with the sensitive microarray-assisted transcriptome profiling. The wild type and a *fixK₂* mutant (strain 9043) were compared as controls. The rationale was that if the Bll2758 protein were a FixT-like negative effector in the wild type, the *bll2758* mutant would display a de-repression of FixK₂ targets. This was clearly not the case (Table 1). Expression of none of the approved FixK₂-activated genes resulted in positive fold-change values in strain 7414 as compared with the wild type (Table 1). Likewise, expression of the *fixK₂* gene itself and of the few other FixJ-P-dependent genes identified previously (Mesa et al. 2008) was unaltered (data not shown). This speaks against a negative regulatory role of Bll2758 in *B. japonicum*, as opposed to the role of FixT-like proteins in *S. meliloti* and *C. crescentus* (Fig. 1).

Negative *fixK₂* autoregulation is possibly mediated by other transcription factors

Lack of evidence for a direct repression of the *fixK₂* gene by the FixK₂ protein led us to assume that autoregulation is indirect. As Bll2758 was excluded here, which other protein might be involved in negative control? A close examination of the comprehensive FixK₂ regulon elaborated by Mesa et al. (2008) uncovered a total of eight genes that code for transcription factors (see box in Fig. 1c). At least one of them might be a candidate to work either as a repressor or as an activator of a repressor gene. The σ^{54} gene *rpoN₁* (*blr1883*) as well as the *fixK₁* and *nnrR* genes (*bll6061* and *blr7084*, respectively) were known to us from previous work (Kullik et al. 1991; Anthamatten et al. 1992; Nellen-Anthamatten et al. 1998; Mesa et al. 2003). Since the σ^{54} and FixK₁ regulons of micro-oxically grown cells were available from genome-wide transcriptomics studies (Hauser et al. 2007; Mesa et al. 2008), it was possible to examine the *fixK₂* gene and all of the 11 direct FixK₂ targets for any de-repression in *rpoN⁻* and *fixK₁⁻* backgrounds, similarly as this was done with the *bll2758⁻* strain (cf. Table 1). Neither *fixK₂* itself nor the 11 targets showed altered expression in the *rpoN⁻* strain N50-97 (data not shown), which argues against them being repressed by a σ^{54} -dependent repressor in the wild-type. Likewise, none of the genes in question was de-repressed in the *fixK₁⁻* strain 7454 (data not shown), arguing against FixK₁ as a potential repressor mediating indirect autoregulation of *fixK₂* in the wild type. To test the NnrR protein as a potential repressor for *fixK₂*, we measured the expression of a chromosomally inserted *fixK₂'-lacZ* fusion in an *nnrR⁻* mutant and, for control, in the wild type and a *fixK₂⁻* mutant (Fig. 6). The *nnrR* mutant showed no difference in *fixK₂*

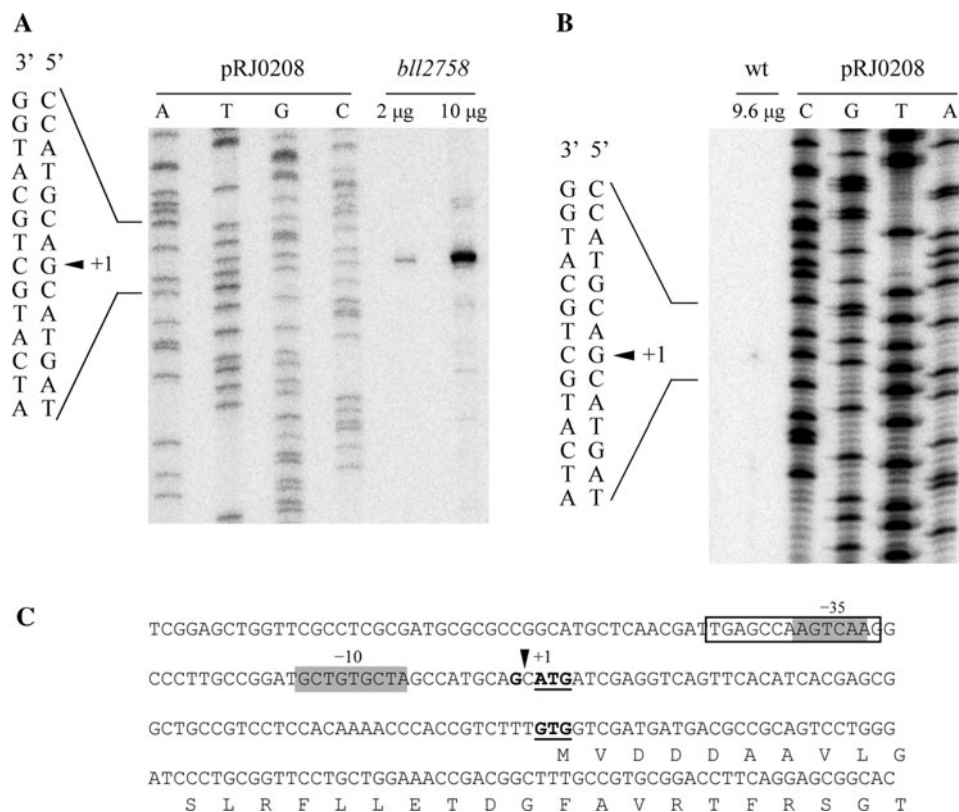


Fig. 4 Transcription start-site determination of *bll2758* by primer extension, and analysis of the promoter region. **a** RNA synthesized by in vitro transcription was used. **b** Total RNA from micro-aerobically grown *B. japonicum* wild-type cells was used. The amounts of RNA applied are indicated above the lanes. Extension products were obtained with the [³²P]-labeled primers 9519-1 (**a**) and LR21 (**b**), and they were separated on a 6% denaturing polyacrylamide gel. The sequencing ladders were generated with plasmid pRJ0208 and primers 9519-1 (**a**) and LR21 (**b**). Part of the promoter region is

shown in the left margin of each panel, with the transcription start site (+1) indicated by an arrowhead. **c** DNA sequence of the *bll2758* promoter region. Again, the transcription start site (+1) is indicated by an arrowhead. The newly proposed (GTG) and the annotated (ATG) translational start codons are underlined. The likely -10 and -35 promoter regions are shaded in gray, and the predicted FixK₂ binding site is boxed. The beginning of the Bll2758 amino acid sequence is indicated in one-letter code

gene expression as compared with the wild type, whereas there was strong *fixK*₂ de-repression in the *fixK*₂ mutant, testifying the negative autoregulation phenomenon. In conclusion, the NnrR protein also does not appear to be involved in the negative control of *fixK*₂.

Discussion

Most of the work reported here deals with the *B. japonicum* *bll2758* gene which codes for a FixT-like single-domain response regulator (Jenal and Galperin 2009). Proteins of that class interfere with other regulators through protein–protein contact instead of binding to critical sites on DNA. A *B. japonicum* *bll2758* null mutant, in contrast to *fixJ* and *fixK*₂ mutants, does not have a conspicuous phenotype with regard to symbiosis and nitrogen fixation (Anthamatten and Hennecke 1991; Nellen-Anthamatten et al. 1998). *A priori*, this does not preclude a negative regulatory role of the

Bll2758 protein in the wild type because, even if knocking the protein out by mutation unleashes its targets, the effect would not necessarily be detectable in the form of a physiologically distinct phenotype. Much more compelling was the microarray experiment done with the *bll2758* mutant, which clearly showed that none of the FixJ- or FixK₂-regulated targets are derepressed.

In conclusion, our work has revealed unexpected species-specific differences in the design of an oxygen-responsive signaling network despite the fact that the regulatory modules (FixL, FixJ, FixT, FixK) are conserved. Particularly puzzling in *B. japonicum* is the FixK₂ dependency of *bll2758* gene expression although its product does not play the role FixT plays in *S. meliloti* or *C. crescentus*. The reason why *bll2758* is activated by FixK₂ remains enigmatic. Yet, negative autoregulation of *fixK*₂ gene expression does exist in *B. japonicum* and, therefore, asks for an alternative mechanistic explanation. While the available evidence suggests an indirect type of control, the

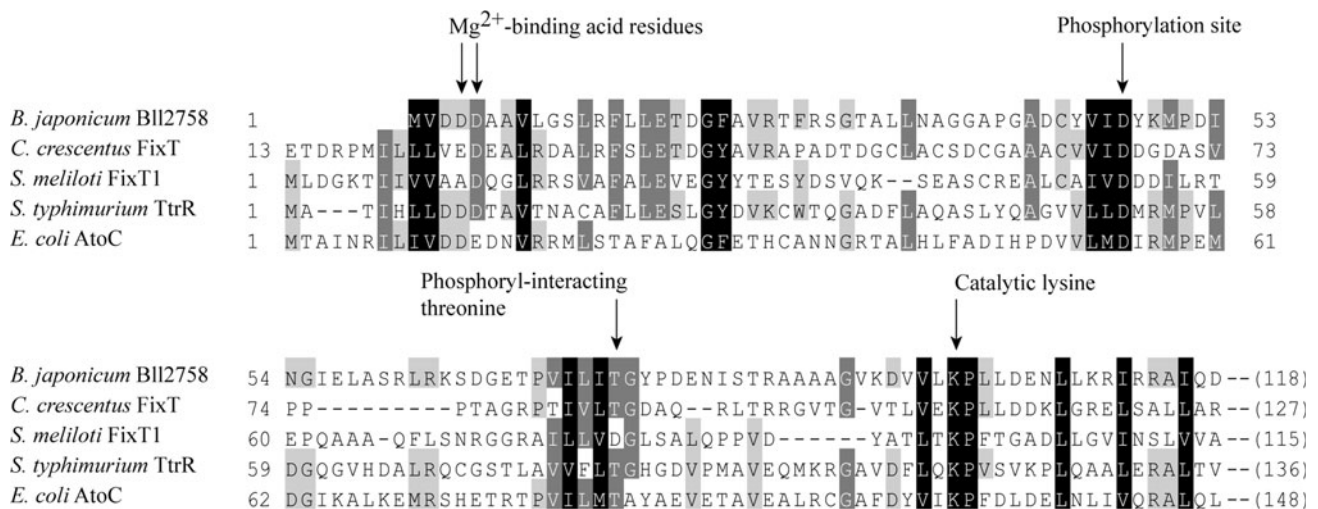


Fig. 5 Alignment of the *B. japonicum* Bll2758 amino acid sequence with other bacterial stand-alone receiver domains. Functionally important amino acid residues of receiver domains are indicated. The protein sequences and their GenBank identification numbers are: *B. japonicum* Bll2758, BAC48023; *C. crescentus* FixT, AAK22738;

S. meliloti FixT1, AAK65326; *S. typhimurium* TtrR, CAB37412; *E. coli* AtoC, AAC75280. While the total number of amino acids of each protein is given at the far end in *parentheses*, a few amino acids from the carboxy-terminal ends are omitted

Table 1 Microarray-assisted expression analysis of the 11 direct FixK₂ targets

Gene number ^b	Gene name	Fold-change of expression compared to wild type ^a	
		Strain 9043 (<i>fixK</i> ₂ ⁻)	Strain 7414 (<i>bll2758</i> ⁻)
<i>blr2763</i>	<i>fixN</i>	-101.9	-
<i>blr2767</i>	<i>fixG</i>	-63.3	-
<i>bll3998</i>		-56.5	-
<i>blr4637</i>	<i>hspC</i> ₂	-111.5	-
<i>blr4655</i>	<i>ppsA</i>	-14.2	-2.3
<i>bll6061</i>	<i>fixK</i> ₁	-19.1	-
<i>blr6062</i>	<i>cycS</i>	-45.9	-
<i>blr6070</i>		-7.3	-
<i>bll6073</i>	<i>phbC</i>	-27.9	-
<i>bll7086</i>	<i>hemN</i> ₂	-97.1	-
<i>bsr7087</i>		-53.8	-

^a All strains were grown under micro-oxic conditions. Negative values indicate decrease of gene expression; - no change within the threshold fold-change range between +2 and -2

^b Gene numbers are from Rhizobase (<http://genome.kazusa.or.jp/rhizobase>). In the case of operons (e.g., *fixNOQP*) only the promoter-proximal gene is listed (i.e., *blr2763*, *fixN*)

possibility cannot be ruled out that FixK₂ represses its own gene directly. If this were the case, however, one would have to postulate a repressor binding motif somewhere around the *fixK*₂ 5' end that radically deviates from the otherwise highly conserved DNA binding site elaborated previously for FixK₂ at activated promoters (Mesa et al. 2005, 2008).

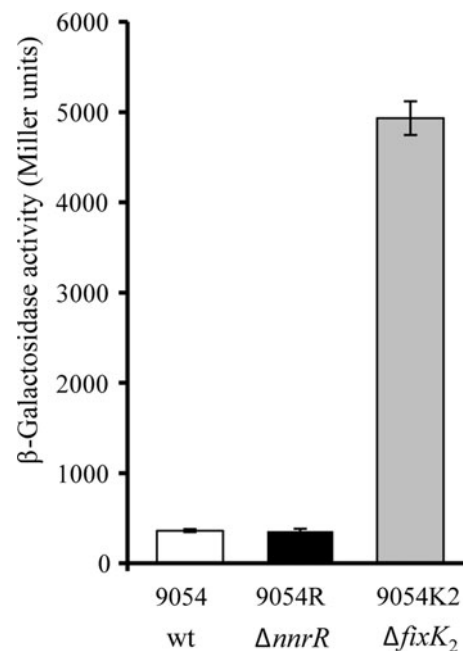


Fig. 6 Expression of a chromosomally integrated *fixK*₂⁻*lacZ* fusion in different *B. japonicum* backgrounds. The relevant genotype of host strains is indicated below strain designations; *wt* wild type. Cells were grown micro-oxically for 72 h before β-galactosidase activity was determined. Values are the means ± standard errors from at least three independent experiments with two cultures assayed in triplicate

Having excluded the Bll2758 protein as a negative effector, we will now have to focus on the other eight FixK₂-dependent genes that encode putative transcription factors. Three of them (*NnrR*, *FixK*₁, *σ*⁵⁴) do not seem to be involved in down-regulating the *fixK*₂ gene. This leaves

not less than five more regulatory genes (*blr1216*, *bll2109*, *bll3466*, *bsr4636*, *blr7666*) for analyses on their potential involvement in autoregulation. To assign such a function to one of them requires a considerable investment into future experimental work. According to the Kazusa annotation (<http://genome.kazusa.or.jp/rhizobase>) *blr1216* codes for a Fur-like repressor, *blr7666* for an AraC-like regulator, and *bsr4636* for a kation transport regulator. The other two genes (*bll2109*, *bll3466*) code for members of the CRP/FNR family of regulators. It will be of interest to find out if any of these regulators acts as a repressor of *fixK₂* or, alternatively, as an activator of a gene that encodes a repressor of *fixK₂*.

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