

RESEARCH PAPER

# Hemoglobin LjGlb1-1 is involved in nodulation and regulates the level of nitric oxide in the *Lotus japonicus*–*Mesorhizobium loti* symbiosis

Mitsutaka Fukudome<sup>1,\*</sup>, Laura Calvo-Begueria<sup>2,\*</sup>, Tomohiro Kado<sup>1</sup>, Ken-ichi Osuki<sup>1</sup>, Maria Carmen Rubio<sup>2</sup>, Ei-ichi Murakami<sup>1,3</sup>, Maki Nagata<sup>1,†</sup>, Ken-ichi Kucho<sup>1</sup>, Niels Sandal<sup>3</sup>, Jens Stougaard<sup>3</sup>, Manuel Becana<sup>2,‡</sup> and Toshiki Uchiumi<sup>1,‡</sup>

<sup>1</sup> Graduate School of Science and Engineering, Kagoshima University, 1-21-35 Korimoto, Kagoshima 890-0065, Japan

<sup>2</sup> Departamento de Nutrición Vegetal, Estación Experimental de Aula Dei, Consejo Superior de Investigaciones Científicas, Apartado 13034, 50080 Zaragoza, Spain

<sup>3</sup> Centre for Carbohydrate Recognition and Signalling, Department of Molecular Biology and Genetics, Aarhus University, Gustav Wieds Vej 10, 8000 Aarhus C, Denmark

\* These authors contributed equally to this work.

† Present address: Institute of Crop Science, National Agriculture and Food Research Organization, 2-1-18 Kannondai, Tsukuba, Ibaraki 305-8518, Japan.

‡ Correspondence: [becana@eead.csic.es](mailto:becana@eead.csic.es) or [uttan@sci.kagoshima-u.ac.jp](mailto:uttan@sci.kagoshima-u.ac.jp)

Received 15 June 2016; Accepted 6 July 2016

Editor: Daphne Goring, University of Toronto

## Abstract

Leghemoglobins transport and deliver O<sub>2</sub> to the symbiosomes inside legume nodules and are essential for nitrogen fixation. However, the roles of other hemoglobins (Hbs) in the rhizobia–legume symbiosis are unclear. Several *Lotus japonicus* mutants affecting LjGlb1-1, a non-symbiotic class 1 Hb, have been used to study the function of this protein in symbiosis. Two TILLING alleles with single amino acid substitutions (A102V and E127K) and a LORE1 null allele with a retrotransposon insertion in the 5′-untranslated region (96642) were selected for phenotyping nodulation. Plants of all three mutant lines showed a decrease in long infection threads and nodules, and an increase in incipient infection threads. About 4h after inoculation, the roots of mutant plants exhibited a greater transient accumulation of nitric oxide (NO) than did the wild-type roots; nevertheless, *in vitro* NO dioxygenase activities of the wild-type, A102V, and E127K proteins were similar, suggesting that the mutated proteins are not fully functional *in vivo*. The expression of LjGlb1-1, but not of the other class 1 Hb of *L. japonicus* (LjGlb1-2), was affected during infection of wild-type roots, further supporting a specific role for LjGlb1-1. In conclusion, the LjGlb1-1 mutants reveal that this protein is required during rhizobial infection and regulates NO levels.

**Key words:** Hemoglobin, *Lotus japonicus*, *Mesorhizobium loti*, nitric oxide, nitrogen fixation, nodulation mutants, symbiosis.

## Introduction

Leghemoglobins are hemoproteins with a relatively high O<sub>2</sub> affinity which are found exclusively in legume nodules (Appleby, 1984; Becana and Klucas, 1992; Smaghe *et al.*, 2009). They transport and deliver O<sub>2</sub> to the symbiosomes at a steady but low concentration to avoid nitrogenase inactivation in bacteroids (Appleby, 1984) and are therefore essential

Abbreviations: Hb, hemoglobin; IT, infection thread; NO, nitric oxide; NOD, nitric oxide dioxygenase; SNAP, S-nitroso-N-acetyl-D,L-penicillamine; SNP, sodium nitroprusside.

© The Author 2016. Published by Oxford University Press on behalf of the Society for Experimental Biology.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/3.0/>), which permits unrestricted reuse, distribution, and reproduction in any medium, provided the original work is properly cited.

for symbiotic N<sub>2</sub> fixation (Ott *et al.*, 2005). However, hemoglobins (Hbs) are not confined to nodules but are ubiquitous in plant tissues, and can be categorized into classes 1, 2, and 3 according to their phylogenetic origin and biochemical properties (Trevaskis *et al.*, 1997; Watts *et al.*, 2001; Hunt *et al.*, 2002; Smaghe *et al.*, 2009).

Class 1 Hbs show extremely high O<sub>2</sub> affinity that disqualifies them as O<sub>2</sub> carriers or sensors (Smaghe *et al.*, 2009). In fact, several lines of evidence indicate that these Hbs are involved in the regulation of intracellular levels of nitric oxide (NO) (Igamberdiev and Hill, 2004; Hebelstrup *et al.*, 2013). This function may be ascribed in part to their NO dioxygenase (NOD) activity: NO+O<sub>2</sub>→NO<sub>3</sub><sup>-</sup> (Igamberdiev and Hill, 2004; Perazzolli *et al.*, 2004). The expression of class 1 Hbs is induced in response to low temperature and hypoxia, conditions known to trigger NO production (Igamberdiev and Hill, 2004; Shimoda *et al.*, 2005). Treatment of plants with NO-releasing compounds, such as sodium nitroprusside (SNP) or *S*-nitroso-*N*-acetyl- D,L-penicillamine (SNAP), also induces expression of class 1 Hb genes (Shimoda *et al.*, 2005; Bustos-Sanmamed *et al.*, 2011). Overexpression of Arabidopsis class 1 Hb (AtGlb1) endows the plants with tolerance to hypoxic stress by decreasing the NO level (Hunt *et al.*, 2002), whereas silencing of the gene gives rise to stunted organs and delayed flowering (Hebelstrup and Jensen, 2008).

The roles of class 1 Hbs may be especially relevant during plant–microbe interactions. In plants, pathogens induce NO production, which in turn activates the expression of pathogenesis-related genes (Delledonne *et al.*, 1998; Durner *et al.*, 1998). NO is rapidly generated in both incompatible and compatible plant–pathogen combinations (Mur *et al.*, 2005), and also participates in disease resistance to necrotrophic pathogens (Asai and Yoshioka, 2009). Ectopic expression of *GhHb1*, a class 1 Hb gene of cotton (*Gossypium hirsutum*), increased the expression of the defense genes *PR1* and *PDF1.2* in Arabidopsis and enhanced resistance to the hemibiotrophic pathogen *Pseudomonas syringae* and the necrotrophic pathogen *Verticillium dahliae* (Qu *et al.*, 2006). On the other hand, an Arabidopsis line with silenced expression of AtGlb1 displayed increased resistance to *P. syringae* and *Botrytis cinerea* (Mur *et al.*, 2012). In this line, the levels of salicylate, jasmonate, and ethylene were increased in response to both pathogens. These findings suggest that class 1 Hbs are involved in the regulation of defense responses through NO and plant hormones. In some, but not all, combinations of symbiotic rhizobia and host plants, inoculation with rhizobia also induces NO generation in the host plant roots, concomitant with the expression of class 1 Hb genes (Nagata *et al.*, 2008). NO inhibits nitrogenase activity (Trinchant and Rigaud, 1982), but is also required for the onset of the *Medicago truncatula*–*Sinorhizobium meliloti* symbiosis (del Giudice *et al.*, 2011). Conversely, the addition of an NO scavenger enhanced the nitrogenase activity of nodules of *Alnus firma* and *Lotus japonicus* (Sasakura *et al.*, 2006; Shimoda *et al.*, 2009). Previous work by Uchiumi and colleagues has shown that overexpression of *LjGlb1-1* (Lj3g3v3338170; [www.kazusa.or.jp/lotus/](http://www.kazusa.or.jp/lotus/)), a class 1 Hb gene of *L. japonicus*,

results in a decrease of NO in nodules, with concomitant increases in nodulation and N<sub>2</sub> fixation of the transformed hairy roots (Shimoda *et al.*, 2009). However, a conclusive proof of the involvement of class 1 Hbs in nodulation is lacking and requires the use of specific mutants because legumes express in nodules several Hbs of all three classes, in addition to leghemoglobins (Uchiumi *et al.*, 2002; Vieweg *et al.*, 2005; Bustos-Sanmamed *et al.*, 2011). In particular, the *L. japonicus* genome contains a second functional class 1 Hb gene, termed *LjGlb1-2* (Lj3g3v3338180; [www.kazusa.or.jp/lotus/](http://www.kazusa.or.jp/lotus/)) (Bustos-Sanmamed *et al.*, 2011).

Here, the symbiotic phenotypes of *L. japonicus* mutant plants that are specifically affected in *LjGlb1-1* expression are reported. This was considered most relevant in the context of NO regulation in the symbiosis because *LjGlb1-1* is the only NO-inducible Hb gene of *L. japonicus* (Bustos-Sanmamed *et al.*, 2011). Two types of mutants, produced by TILLING (Perry *et al.*, 2009) and insertion of retrotransposon LORE1 (Fukai *et al.*, 2012; Urbański *et al.*, 2012; Małolepszy *et al.*, 2016), were used in this work and shown to have lower infection rates, fewer nodules, and a higher NO level in roots than the wild-type (WT) plants. We conclude that *LjGlb1-1* supports *M. loti* infection, probably by regulating the NO level in the roots of *L. japonicus*.

## Materials and methods

### Biological material and plant growth

Plants of WT and mutant lines of *L. japonicus* accession Gifu B-129 were used in this study. Seeds were gently scarified, disinfected with 2% sodium hypochlorite, washed, and left overnight in darkness. After imbibition, seeds were washed, left on plates with 0.5% agar for 3 d at 4 °C, transferred to agar plates containing nutritive medium, and placed vertically at 24 °C for 3 d in the dark. For phenotype analyses, seedlings were grown on Fåhraeus (1.5% agar) medium (Fåhraeus, 1957) and each seedling was inoculated with 10<sup>6</sup> cells of *M. loti* strains MAFF303099 or MAFF303099 *DsRed* (Maekawa *et al.*, 2009). For quantitative real-time PCR (qPCR) analyses, seedlings were grown on Jensen (0.9% agar) medium (Pajuelo and Stougaard, 2005) and inoculated with 10<sup>7</sup> cells of *M. loti* strain R7A. Roots were harvested 1, 2, 4, and 6 d after inoculation, and were immediately flash-frozen in liquid nitrogen and stored at –80 °C until use. Uninoculated roots of the same age served as controls. In plates, roots were protected from light using black cardboard. For phenotyping non-nodulated plants, the Fåhraeus medium was supplemented with 1.5 mM NH<sub>4</sub>NO<sub>3</sub>. Plates were placed in growth cabinets with a 24 °C/21 °C day/night regime, 16 h photoperiod, and 150 μmol m<sup>-2</sup> s<sup>-1</sup> light intensity.

### Mutant lines from TILLING and LORE1 populations

TILLING was performed by RevGenUK (<http://revgenuk.jic.ac.uk/>), which provided M<sub>3</sub> seeds carrying mutations in *LjGlb1-1*. Genomic DNA was isolated from leaves of M<sub>4</sub> progeny with a DNeasy Plant Kit (Qiagen) according to the manufacturer's instructions. Homozygous plants were selected by PCR with primers 5'-AGT CTA GAG TAA TCA CAT CAA TTC CAC C-3' and 5'-TGA GTC TAA GAA GAT GAT GGC TTC A-3' using a program consisting of 25 cycles at 94 °C for 30 s, 57 °C for 1 min, and 72 °C for 1 min. The amplified products were sequenced. The M<sub>5</sub> progeny derived from each homozygous individual were used for further experiments.

Null mutants bearing a retrotransposon insertion in the 5'-untranslated region (UTR) (line 30096642, hereafter abbreviated as 96642

to simplify) were obtained from the *LORE1* collection (Fukai *et al.*, 2012; Urbański *et al.*, 2012; Małolepszy *et al.*, 2016). The plants in the segregating population were genotyped by PCR using 96642-fw primer 5'-CAT GGC ATG AGG CTT GAG CTT GGG-3' and 96642-rev primer 5'-TGA AAC CAC TCT CTT CTC GCC GCA-3' to amplify the WT copy of *LjGlb1-1* and with 96642-fw primer and *LORE1* primer P2 5'-CCA TGG CGG TTC CGT GAA TCT TAG G-3' to amplify the *LjGlb1-1* copy with the *LORE1* insertion. Three individual homozygous mutant plants were selected for analysis. Because their growth phenotypes did not significantly differ, the data obtained with the three individual mutants were pooled in this study. Both *TILLING* and *LORE1* mutant alleles originate from the *L. japonicus* Gifu ecotype (Handberg and Stougaard, 1992).

#### Nodulation and nitrogenase activity

Plants grown as indicated earlier on Fåhræus medium were inoculated with  $10^6$  cells of *M. loti* MAFF303099 or its *DsRed*-tagged derivative. Two weeks after inoculation, infection threads (ITs) were counted in two groups (incipient and long ITs) according to the terminology of Małolepszy *et al.* (2015), except that in our case the elongating ITs were also considered as long ITs. Four weeks after inoculation, nodules were detached and counted. The size and fresh weight of each plant were also measured. Nitrogenase activity of the detached nodules was determined as acetylene reduction activity according to Shimoda *et al.* (2009).

#### Microscopic observation of endogenous and released NO in roots

Endogenous NO generation in roots of 5-day-old seedlings was monitored by fluorescence microscopy as described (Nagata *et al.*, 2008). Seedlings were inoculated with *M. loti* MAFF303099 or mock treated (sterile distilled water) and incubated for 3 h. The seedlings were then soaked for 1 h with 20  $\mu$ M 4-amino-5-methylamino-2',7'-difluorescein diacetate (DAF-FM DA; Sekisui Medical, Japan). In some experiments, the NO scavenger 2-(4-carboxyphenyl)-4,4,5,5-tetramethylimidazoline-1-oxyl-3-oxide (cPTIO) was applied at a concentration of 3 mM simultaneously with DAF-FM DA. Confocal images were captured with an A1si-90i microscope (Nikon, Japan) and epifluorescence images with an Eclipse 90i microscope (Nikon, Japan).

The concentration of NO released from the roots was assessed 4 h after inoculation with *M. loti* MAFF303099. Roots were incubated with 7  $\mu$ M DAF-FM for 3 min, and the fluorescence intensity of the DAF-FM solution was measured as described (Tominaga *et al.*, 2009) with an ES-2 micro UV-visible fluorescence spectrophotometer (Malcom; Japan) using 495 nm and 519 nm as excitation and emission wavelengths, respectively. The released NO was expressed as relative fluorescence intensity per fresh weight of roots.

To investigate the effect of exogenous NO, 3-day-old seedlings were inoculated with *M. loti* MAFF303099 *DsRed*, and 500  $\mu$ M SNAP or 100  $\mu$ M SNP was applied to the roots alone or with 500  $\mu$ M cPTIO at the same time as inoculation. Two weeks after inoculation, the numbers of incipient and long ITs were counted.

#### Production and mutagenesis of recombinant LjGlb1-1

The LjGlb1-1 protein (WT) was expressed in Champion pET200/D-TOPO vector (Invitrogen) as previously described in detail (Sainz *et al.*, 2013). The mutated derivative E127K was generated by PCR-based substitution by Mutagenex (Somerset, NJ, USA). Both the DNA and protein were sequenced to confirm the amino acid substitution. The proteins bearing an N-terminal poly-His tag were expressed in *Escherichia coli* C41 (DE3) cells (Lucigen) at 37 °C for 4–6 h with 0.2 mM isopropyl- $\beta$ -D-l-thiogalactopyranoside. Cells were resuspended in 50 mM potassium phosphate (pH 7.5), broken by sonication, and cleared by centrifugation. The supernatant was fractionated with 30–75% ammonium sulfate, dialyzed in phosphate-buffered saline [PBS; 50 mM potassium phosphate (pH

7.5)+150 mM NaCl], and loaded on an Ni affinity column (HiTrap Chelating HP; GE Healthcare). The column was washed with 5 vols of PBS+20 mM imidazole, and the recombinant proteins were eluted with PBS+250 mM imidazole. The proteins were oxidized with ferricyanide, dialyzed in PBS, concentrated, quantified based on the Soret absorption band, and stored at –80 °C until analysis.

#### Spectra and NOD activities of recombinant LjGlb1-1 proteins

The Soret–visible spectra of the LjGlb1-1 proteins (WT and E127K) in the ferric ( $\text{Hb}^{3+}$ ), deoxyferrous ( $\text{Hb}^{2+}$ ), and oxyferrous ( $\text{Hb}^{2+}\text{O}_2$ ) states were recorded with a UV-visible Lambda 25 spectrophotometer (Perkin-Elmer).  $\text{Hb}^{2+}$  was produced by adding a trace of dithionite to  $\text{Hb}^{3+}$ , and  $\text{Hb}^{2+}\text{O}_2$  was formed by passing  $\text{Hb}^{2+}$  through a NAP-5 mini-column (GE Healthcare).

The NOD activities of the recombinant proteins were measured by using two NO donors and an NO-specific electrode (ISO-NOP), which was calibrated daily following the manufacturer's instructions (World Precision Instruments, Sarasota, FL, USA). Diethylamine NONOate (DEA; 20  $\mu$ M) and *S*-nitrosoglutathione (GSNO; 1 mM) were added to 4 ml of 50 mM potassium phosphate buffer (pH 7.5) containing 50  $\mu$ M diethylenetriaminepentaacetic acid. This solution was kept with gentle shaking at 24 °C until the NO concentration became stable (~4 min), and freshly prepared  $\text{Hb}^{2+}\text{O}_2$  protein (2  $\mu$ M) was added and the decrease of NO measured. The time between the preparation of  $\text{Hb}^{2+}\text{O}_2$  proteins and the NOD activity assay was always <5 min. The  $\text{Hb}^{3+}$  proteins lacked NOD activity and were used as controls.

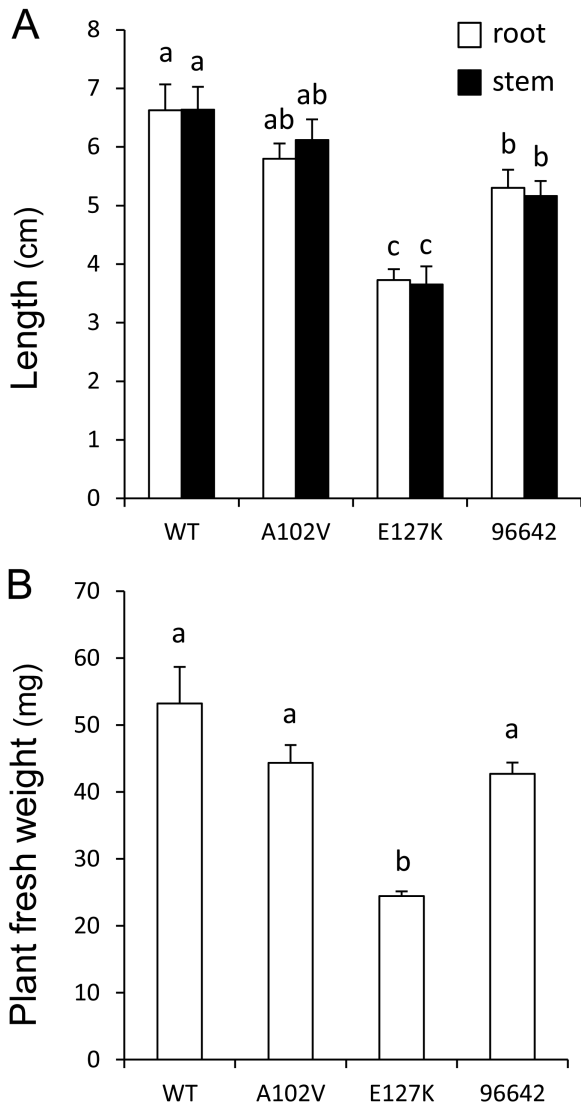
#### Real-time quantitative PCR

Total RNA was extracted from 40–70 mg of roots (6–10 roots) with the RNAqueous isolation kit (Ambion) and cDNA was synthesized using DNase-treated RNA with (dT)<sub>17</sub> and Moloney murine leukemia virus reverse transcriptase (Promega). qPCR analysis was performed using a 7500 Real-Time PCR system (Applied Biosystems) and iTaq Universal SYBR Green Supermix reagents (Bio-Rad). Primers for *LjGlb1-1* (5'-TCT CAC TTC ACT TCC ATC GCA-3' and 5'-TCA GTG AAA CAT GTG CTC CCA-3') and *LjGlb1-2* (5'-GGC AGA AAA CAC AAC CAC CAT-3' and 5'-TCA CCA CCA GAG CTT CTT GCT-3') were used with a PCR program consisting of an initial denaturation and *Taq* polymerase activation step of 10 min at 95 °C, followed by 40 cycles of 15 s at 95 °C and 1 min at 60 °C, and a final melting curve stage. Primer specificity and the absence of contaminating genomic DNA were verified, respectively, by amplicon dissociation curves and by PCR analysis of RNA samples prior to reverse transcription. Expression levels were normalized using *ubiquitin* as reference gene, which was found to remain constant in roots during the few days of measurements.

## Results

### Growth and nodulation phenotypes of mutant LjGlb1-1 lines

In this work, two types of LjGlb1-1 mutants were used. On the one hand, mutant heterozygotes were identified by *TILLING*, each bearing a different amino acid substitution. Four mutant homozygous lines were established by self-pollination and designated as P59S, A102V, E127K, and E136K according to the amino acid substitutions and their positions. All of them showed a lower plant length than the WT, but only the A102V and E127K lines were selected for further studies because A102 is conserved and close to H104, which is important for assembling the heme moiety (Andersson *et al.*, 1996), and because E127 is highly conserved among



**Fig. 1.** Growth phenotype of *LjGlb1-1* mutant lines. Three days after germination, seedlings were inoculated with *M. loti* MAFF303099 and grown on nitrogen-free Fåhræus medium for 4 weeks. At this time, (A) the root and stem lengths and (B) the fresh weight of plants were measured. Means ( $\pm$  SE;  $n=18$  for WT,  $n=30$  for 96642,  $n=12-16$  for the other lines) denoted by the same letter do not significantly differ based on Duncan's multiple range test at  $P=0.05$ .

**Table 1.** Nodulation phenotype of *LjGlb1-1* mutant lines

Seedlings were inoculated with *M. loti* MAFF303099 (nodule number and nitrogenase activity) or its *DsRed* derivative (ITs) and were grown on nitrogen-free Fåhræus medium. Nitrogenase (acetylene reduction) activity (ARA) of detached nodules was measured after 4 weeks and is expressed as  $\text{nmol C}_2\text{H}_4 \text{ produced min}^{-1} \text{ mg}^{-1} \text{ FW}$ . The numbers of ITs were counted after 2 weeks and are expressed per centimeter of root.

	WT	A102V	E127K	96642
Nodule number	7.17 $\pm$ 0.51 a	5.22 $\pm$ 0.30 b	2.67 $\pm$ 0.28 c	4.93 $\pm$ 0.42 b
ARA	11.18 $\pm$ 0.72 a	9.64 $\pm$ 0.51 ab	2.71 $\pm$ 0.60 c	8.54 $\pm$ 0.69 b
Incipient ITs	2.88 $\pm$ 0.13 a	4.43 $\pm$ 0.42 b	8.15 $\pm$ 1.28 b	4.52 $\pm$ 0.61 b
Long ITs	32.08 $\pm$ 1.41 a	17.11 $\pm$ 0.98 b	5.98 $\pm$ 0.93 c	11.57 $\pm$ 0.82 d
Total ITs	34.95 $\pm$ 1.42 a	21.54 $\pm$ 1.30 b	14.00 $\pm$ 2.05 c	16.09 $\pm$ 1.21 c

Means ( $\pm$  SE,  $n=9$  for ARA and  $n=14-22$  for the other parameters) denoted by the same letter do not significantly differ ( $P=0.05$ ) based on Duncan's multiple range test.

class 1 plant Hbs (Supplementary Fig. S1 at *JXB* online). On the other hand, the mutant line 96642, which contains an insertion in the 5'-UTR of the *LjGlb1-1* gene 11 bp upstream of ATG, was selected from the *LORE1* collection. Lines with annotated insertions in the coding parts of the exons of *LjGlb1-1* were not found in a total of 134 682 lines. Seedlings of the 96642 line were grown on agar plates and homozygous mutant plants were transferred to the greenhouse for seed production. Detailed qPCR analyses revealed that the *LjGlb1-1* mRNA level in roots of the A102V and E127K lines is similar to that of the WT, whereas it was virtually undetectable in the 96642 line ( $<1\%$  of WT). This percentage was calculated considering the threshold value ( $C_t$ ) for *LjGlb1-1* in the 96642 (32–34 cycles) and WT (25–31 cycles) plants, together with the  $C_t$  value for *ubiquitin* (16–19 cycles). Consequently, the 96642 line can be considered as a null mutant.

The two TILLING and the single *LORE1* mutant lines were used for phenotyping, which included measurements of lengths and fresh weights of 4-week-old plants nodulated with *M. loti* MAFF303099. The E127K and 96642 plants had shorter roots and stems (Fig. 1A) and the E127K plants weighed less (Fig. 1B) than the WT. However, all the mutants maintained the same shoot length/root length ratio as the WT (Fig. 1A). The three mutant lines formed fewer nodules and the E127K and 96642 lines exhibited significantly lower  $\text{N}_2$  fixation activity, estimated as acetylene reduction activity per fresh weight of nodules (Table 1). The roots of the WT and mutant plants were inoculated with the *DsRed*-labeled strain and the numbers of incipient ITs (microcolonies+short ITs) and long ITs (Fig. 2) were counted 2 weeks after inoculation. Compared with the WT plants, the numbers of incipient ITs were increased in the three mutant lines, whereas the numbers of long and total ITs were reduced (Table 1). Taken together, these observations indicate an alteration in the progression of ITs during infection in the roots of mutant plants.

Two additional phenotype measurements were carried out. First, the w/w siblings derived from  $M_4$  seeds of the A102V and E127K plants did not show significant differences in the plant length and in the nodule and IT numbers (Supplementary Table S1), which suggests that the mutations in the *LjGlb1-1* gene are responsible for the observed phenotypes. Secondly,

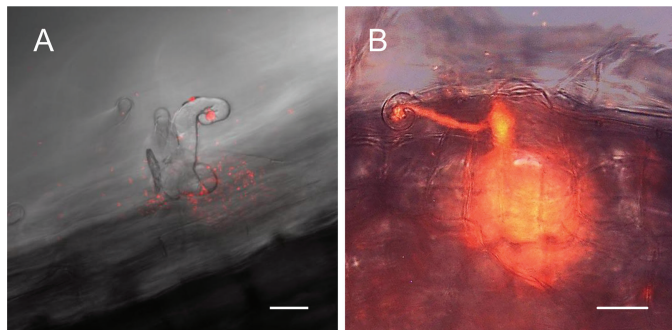
because *LjGlb1-1* is expressed in roots and leaves in addition to nodules (Nagata *et al.*, 2008; Bustos-Sanmamed *et al.*, 2011), the phenotype was also analyzed in non-nodulated plants fed with  $\text{NH}_4\text{NO}_3$  (Supplementary Table S2). Except for the shoot weight and length of the A102V plants, the growth parameters of the three mutant lines were decreased with respect to the WT. These results indicate that LjGlb1-1 does not only affect nodulation but is also involved in plant growth and development, as would be expected for a protein that is highly, but non-specifically, expressed in the nodules.

#### Involvement of NO in the *LjGlb1-1* nodulation phenotype

The NO levels in the roots of WT and mutant plants were compared because class 1 Hbs may modulate the NO concentration *in vivo* (Hebelstrup *et al.*, 2013; Wally *et al.*, 2013). For this purpose, the NO-specific fluorescent dye DAF-FM DA was applied to the roots 3 h after inoculation with *M. loti* and then incubated for 1 h. This compound is cell permeant and essentially non-fluorescent, but it is deacetylated by intracellular esterases to DAF-FM, which in turn reacts with endogenous NO forming a highly fluorescent benzotriazole. In the WT roots, the fluorescence intensity, marking the production of NO, increased very slightly 4 h after inoculation with *M. loti* (Fig. 3), consistent with a previous report (Nagata

*et al.*, 2008). However, the signal intensity was enhanced in the A102V, E127K, and 96642 mutants compared with the WT (Fig. 3). This signal was abolished by incubation with the NO-specific scavenger cPTIO, thus confirming that NO was the reactive molecule being detected (Supplementary Fig. S2). Consequently, endogenous NO was low and increased slightly in the WT roots shortly after infection, but this transient increase was higher in the mutant roots. The NO released from roots was quantified by using the non-permeant dye DAF-FM and expressed as the relative fluorescence signal per fresh weight of roots. The fluorescence intensity was significantly higher in the roots of the three mutant plants than in the roots of WT plants (Fig. 4), in agreement with the observed increase of NO production by the roots of both mutants lines.

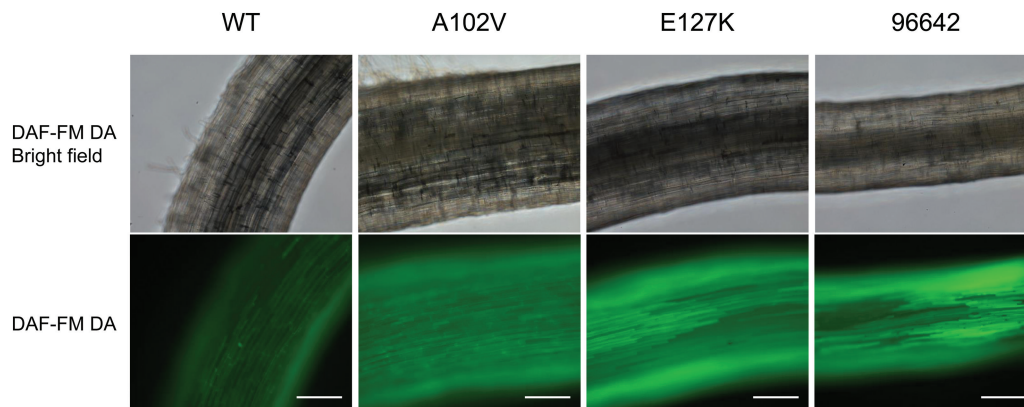
To investigate further whether NO was mediating, at least in part, the nodulation phenotype of the mutants, SNAP was applied to roots of WT plants at the same time as inoculation. Treatment with this NO donor resulted in an increase in the number of incipient ITs and in a reduction of the numbers of long and total ITs (Table 2). The addition of cPTIO alone or together with SNAP reverted completely or partially the effect of SNAP on those parameters, pointing to NO as the responsible reactive molecule (Table 2). Similar increasing (incipient ITs) or decreasing (long ITs) effects were observed upon the treatment of roots with SNP, another NO donor (Supplementary Table S3).



**Fig. 2.** Typical (A) incipient IT and (B) long IT in roots of WT plants after 14 d of inoculation. Images were taken with a confocal microscope. Scale bars=25  $\mu\text{m}$ .

#### Characterization of mutated *LjGlb1-1*

The possibility that the increased NO accumulation observed in roots of A102V and E127K plants was due to an alteration of the biochemical properties of the mutated proteins was explored. To this end, recombinant WT, A102V, and E127K proteins were purified by metal-affinity chromatography. The preparations had a purity of >90% and were free of contaminating *E. coli* flavohemoglobin (44 kDa) according to Coomassie-stained SDS gels (Supplementary Fig. S3). The absence of contaminating Hb from *E. coli* was further confirmed using the same vector and purification protocol from bacterial extracts for glutathione peroxidase, which

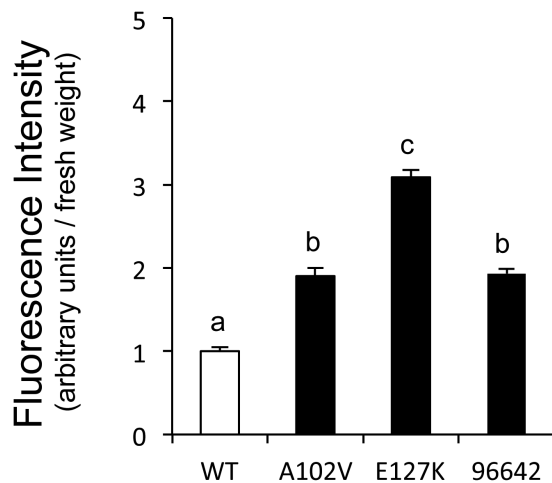


**Fig. 3.** Fluorescence imaging of NO production in the roots of WT and mutant lines. Seedlings were incubated with *M. loti* MAFF303099 for 3 h and then with DAF-FM DA for 1 h. The images of the roots for all three lines were taken with a confocal microscope using identical settings. Roots incubated with mock (sterile distilled water) instead of rhizobia did not show detectable fluorescence. Scale bars=200  $\mu\text{m}$ .

lacked NOD activity as expected for a non-heme protein. The Soret-visible spectra (Supplementary Fig. S4) and NOD activities (Fig. 5) of the three proteins were compared. The spectra were similar, indicating that the hemes in the mutated proteins are hexacoordinated in the ferric and deoxyferrous forms (Smaghe et al., 2009; Sainz et al., 2013) and are still able to bind O<sub>2</sub>. This was confirmed by the ability of the A102V and E127K proteins to carry out the NOD reaction, which requires both O<sub>2</sub> and NO binding inside the heme cavity. In fact, the mutated proteins displayed *in vitro* similar NOD activities to the WT protein using DEA or GSNO as NO donors (Fig. 5). Consequently, the potential of the mutant proteins to scavenge NO through its dioxygenation to NO<sub>3</sub><sup>-</sup> remains intact.

#### Expression profiles of LjGlb1-1 and LjGlb1-2 during infection

The *L. japonicus* genome encodes two class 1 Hbs, but only LjGlb1-1 appears to be induced by NO at the transcriptional level (Shimoda et al., 2005; Bustos-Sanmamed et al., 2011). It was therefore of interest to determine by qPCR the expression



**Fig. 4.** Quantification of NO released from roots of WT and mutant plants after incubation with *M. loti* MAFF303099. The fluorescence intensity of the DAF-FM solution in the rooting medium was measured with a plate reader and expressed as relative fluorescence intensity per fresh weight of roots. Means ( $\pm$  SE;  $n=9$ ) denoted by the same letter do not significantly differ based on Duncan's multiple range test at  $P=0.05$ .

**Table 2.** Effect of NO on nodulation of WT plants

Seedlings were inoculated with *M. loti* MAFF303099 DsRed and 100  $\mu$ l of 500  $\mu$ M SNAP, 500  $\mu$ M cPTIO, or a combination of both compounds, was applied on each root. The numbers of ITs are expressed per centimeter of root. All parameters were measured 2 weeks after inoculation.

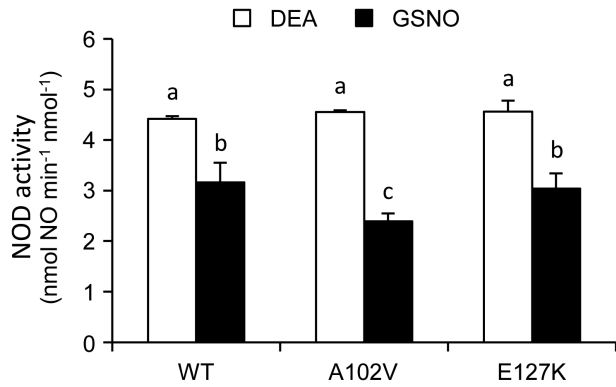
	Control	SNAP	cPTIO	SNAP+cPTIO
Incipient ITs	2.7 $\pm$ 0.3 a	8.0 $\pm$ 1.1 b	4.9 $\pm$ 0.6 b	3.8 $\pm$ 0.5 a
Long ITs	25.4 $\pm$ 2.7 a	9.8 $\pm$ 1.2 b	28.7 $\pm$ 2.4 a	26.0 $\pm$ 3.3 a
Total ITs	28.1 $\pm$ 3.0 a	17.8 $\pm$ 1.9 b	33.6 $\pm$ 2.4 a	29.7 $\pm$ 3.7 a

Means ( $\pm$  SE,  $n=9-11$ ) denoted by the same letter do not significantly differ ( $P=0.05$ ) based on Duncan's multiple range test.

profiles of the *LjGlb1-1* and *LjGlb1-2* genes in roots of WT plants during the first days of infection (Fig. 6). The expression profile of *LjGlb1-2* was also obtained for roots of the 96642 line, which shows undetectable *LjGlb1-1* mRNA levels. In these experiments, uninoculated roots of the same age were used for comparison (Fig. 6, white bars). The *LjGlb1-1* mRNA level of inoculated roots decreased between 1d and 6 d after infection (Fig. 6A, black bars), whereas the *LjGlb1-2* mRNA level was not affected (Fig. 6B, black bars). The transcript levels of both genes remained constant in the uninoculated root controls (Fig. 6A, B, white bars). No changes were observed for the *LjGlb1-2* mRNA levels in the roots of the 96642 mutant (Fig. 6C, white and black bars).

## Discussion

Previous work in one of our laboratories using transgenic hairy roots of *L. japonicus* has shown that overexpression of *LjGlb1-1* improves nodule formation, decreases the NO level, and enhances nitrogenase activity of nodules (Shimoda et al., 2009). These observations led us to surmise that a deficiency of LjGlb1-1 in plants would reduce nodulation and nitrogenase activity. To test this hypothesis, three mutant lines of *L. japonicus* were used in this study. The A102V and E127K lines express proteins bearing single mutations in conserved amino acid residues, whereas the 96642 line is unable to express the protein. The E127K and 96642 plants showed shorter roots and stems and the E127K plants had lower fresh weight (Fig. 1). The three mutants were also altered in nodulation, with decreases of the nodule number, long ITs, and acetylene reduction activity, and an increase of incipient ITs (Table 1). However, the plant growth and nodulation were more inhibited in E127K than in A102V and 96642, whereas w/w siblings of both A102V and E127K plants did not show this phenotype (Supplementary Table S1). A similar more intense phenotype was observed for E127K plants grown under non-nodulating conditions, which also disappears in the w/w siblings (Supplementary Table S2). Thus, it may be that the E127K line carries an additional linked mutation causing, at least in part, a more penetrating phenotype. The reduction in the number of long ITs in the roots of the three mutant lines indicates that the infection process is restricted at the initial stages of rhizobial entry, particularly during progression of ITs. This alteration may, in turn, increase in

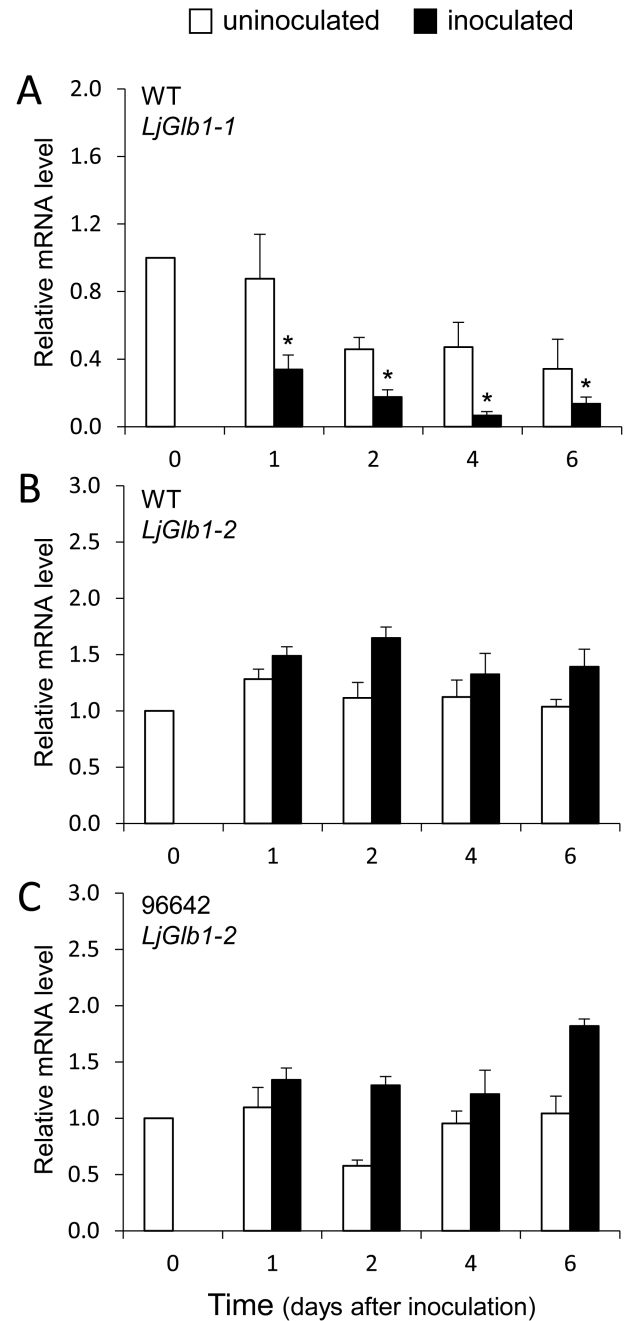


**Fig. 5.** NOD activity of recombinant LjGlb1-1 WT and its A102V and E127K mutant derivatives. The activity was measured with 20  $\mu$ M DEA or 1 mM GSNO and 2  $\mu$ M oxyferrous proteins in 50 mM potassium phosphate buffer (pH 7.5) containing 50  $\mu$ M diethylenetriaminepentaacetic acid. Means ( $\pm$  SE;  $n=2-3$  independent protein preparations) denoted by the same letter do not significantly differ based on Duncan's multiple range test at  $P=0.05$ .

the number of attempted infections, as observed for other mutants (Wopereis *et al.*, 2000).

The results of this work also reveal that this negative effect on nodulation is due, at least in part, to deregulation of NO levels in the roots. Thus, NO accumulates in roots after only ~4 h from infection, confirming our previous observation (Shimoda *et al.*, 2005). Here, the accumulation of NO was found to be enhanced in the roots of the three mutant lines, providing strong support for our hypothesis that LjGlb1-1 regulates endogenous NO levels during infection so as to avoid triggering of a defense response in roots (Shimoda *et al.*, 2005). The transient increase of endogenous NO would therefore explain the alteration of the infection process, as evidenced by the changes in the nodule and IT numbers in the mutant plants (Table 1). This proposal is further supported by our observation that exogenous NO, provided by SNAP or SNP, also inhibits IT elongation (Table 2). The reason why the roots of the A102V and E127K plants accumulate higher NO levels than those of the WT plants is nevertheless uncertain because the two mutated proteins and the WT protein exhibit similar NOD activities *in vitro* (Fig. 5). Two plausible and compatible explanations are that the NOD activity is being limited *in vivo* by a reductant that is essential to regenerate the ferrous globin (Becana and Klucas, 1992; Igamberdiev and Hill, 2004; Smaghe *et al.*, 2008; Sainz *et al.*, 2013), and that the mutations affect protein stability in the intracellular medium. An alternative explanation that cannot be discarded, but seems more unlikely, is that the higher accumulation of NO in the mutants is indirectly originated by LjGlb1-1 malfunction rather than by a decreased capacity for NO scavenging.

The expression analysis of the two class 1 Hb genes of *L. japonicus* shows that *LjGlb1-1* mRNA levels decrease within the first days of infection (Fig. 6A). Previous work has shown that *LjGlb1-1* is induced by NO after only 3–4 h (Shimoda *et al.*, 2005; Bustos-Sanmamed *et al.*, 2011), and therefore NO and *LjGlb1-1* expression levels might change in parallel during infection. In this scenario, NO accumulation



**Fig. 6.** Expression analysis of the *LjGlb1-1* and *LjGlb1-2* genes in roots during the first days upon inoculation with *M. loti* R7A. Transcript steady-state levels were expressed relative to the day of inoculation (day 0). The figure shows expression levels of *LjGlb1-1* (A) and *LjGlb1-2* (B) in roots of WT plants, and of *LjGlb1-2* (C) in roots of 96642 plants. Values are means  $\pm$  SE of four biological replicates (different RNA extractions), each with three technical replicates. White and black bars represent values for uninoculated and inoculated roots, respectively. The asterisk denotes significant gene down-regulation (relative mRNA level  $<0.5$ ).

and hence *LjGlb1-1* induction are restricted to a short time frame, and afterwards, between 1 d and 6 d, both NO and *LjGlb1-1* mRNA levels decline (Shimoda *et al.*, 2005; Fig. 6A). In sharp contrast, *LjGlb1-2* expression remained unaffected (Fig 6B, C), which allows us to conclude that this gene is not involved in the NO-dependent response of the roots to rhizobial infection.

In the *M. truncatula*–*S. meliloti* interaction, NO is required for the optimal establishment of the symbiosis, and a decrease of NO production in the roots inhibits nodulation (del Giudice et al., 2011; Pauly et al., 2011). Our results, including those of Shimoda et al. (2005) and Nagata et al. (2008), appear to contradict these reports, and so far there is no clear explanation for this discrepancy, which may be ascribed to differences between the indeterminate (*M. truncatula*) and determinate (*L. japonicus*) patterns of nodule development and/or in the range of NO concentrations needed for the onset of the symbioses. In any case, the use of LjGlb1-1 mutants in the present study provides conclusive evidence that this particular class 1 Hb is essential for proper nodulation as its deficiency results in a reduced number of elongated ITs caused, at least in part, by the transient accumulation of NO in the infected roots.

## Supplementary data

Supplementary data are available at *JXB* online.

**Table S1.** Growth and nodulation phenotype of WT plants and w/w siblings ( $M_3$  seeds) from mutant plants.

**Table S2.** Growth parameters of non-nodulated LjGlb1-1 mutant plants, and derived w/w siblings, supplied with combined nitrogen.

**Table S3.** Effect of SNP application to roots on nodulation of WT plants.

**Figure S1.** Alignment of some class 1 Hbs showing conservation of A102 and E127.

**Figure S2.** Inhibition of NO-associated fluorescence by cPTIO in roots of WT and mutant plants.

**Figure S3.** Purification of WT recombinant LjGlb1-1.

**Figure S4.** Representative Soret–visible spectra of recombinant WT, A102V, and E127K proteins.

## Acknowledgements

We thank the National BioResource Project for providing seeds of *L. japonicus* Gifu B-129, and Carmen Pérez-Rontomé for help with the NOD activity assays. This project was supported by Open Partnership Joint Projects of JSPS Bilateral Joint Research Projects (Japan), by Danish National Research Foundation grant DNRF79 (Denmark), and by Ministry of Economy and Competitiveness grant AGL2014-53717-R (Spain).

## References

- Andersson CR, Jensen EO, Llewellyn DJ, Dennis ES, Peacock WJ. 1996. A new hemoglobin gene from soybean: a role for hemoglobin in all plants. *Proceedings of the National Academy of Sciences, USA* **93**, 5682–5687.
- Appleby CA. 1984. Leghemoglobin and *Rhizobium* respiration. *Annual Review of Plant Physiology* **35**, 443–478.
- Asai S, Yoshioka H. 2009. Nitric oxide as a partner of reactive oxygen species participates in disease resistance to necrotic pathogen *Botrytis cinerea* in *Nicotiana benthamiana*. *Molecular Plant-Microbe Interactions* **22**, 619–629.
- Becana M, Klucas RV. 1992. Oxidation and reduction of leghemoglobin in root nodules of leguminous plants. *Plant Physiology* **98**, 1217–1221.
- Bustos-Sanmamed P, Tovar-Méndez A, Crespi M, Sato S, Tabata S, Becana M. 2011. Regulation of nonsymbiotic and truncated hemoglobin

genes of *Lotus japonicus* in plant organs and in response to nitric oxide and hormones. *New Phytologist* **189**, 765–776.

del Giudice J, Cam Y, Damiani I, Fung-Chat F, Meilhoc E, Bruand C, Brouquisse R, Puppo A, Boscari A. 2011. Nitric oxide is required for an optimal establishment of the *Medicago truncatula*–*Sinorhizobium meliloti* symbiosis. *New Phytologist* **191**, 405–417.

Delledonne M, Xia YJ, Dixon RA, Lamb C. 1998. Nitric oxide functions as a signal in plant disease resistance. *Nature* **394**, 585–588.

Durner J, Wendehenne D, Klessig DF. 1998. Defense gene induction in tobacco by nitric oxide, cyclic GMP, and cyclic ADP-ribose. *Proceedings of the National Academy of Sciences, USA* **95**, 10328–10333.

Fähræus G. 1957. The infection of clover root hairs by nodule bacteria studied by a single glass slide technique. *Microbiology* **16**, 374–381.

Fukai E, Soyano T, Umehara Y, Nakayama S, Hirakawa H, Tabata S, Sato S, Hayashi M. 2012. Establishment of a *Lotus japonicus* gene tagging population using the exon-targeting endogenous retrotransposon LORE1. *The Plant Journal* **69**, 720–730.

Handberg K, Stougaard J. 1992. *Lotus japonicus*, an autogamous, diploid legume species for classical and molecular genetics. *The Plant Journal* **2**, 487–496.

Hebelstrup KH, Jensen HO. 2008. Expression of NO scavenging hemoglobin is involved in the timing of bolting in *Arabidopsis thaliana*. *Planta* **227**, 917–927.

Hebelstrup KH, Shah JK, Igamberdiev AU. 2013. The role of nitric oxide and hemoglobin in plant development and morphogenesis. *Physiologia Plantarum* **148**, 457–469.

Hunt PW, Klok EJ, Trevaskis B, Watts RA, Ellis MH, Peacock WJ, Dennis ES. 2002. Increased level of hemoglobin 1 enhances survival of hypoxic stress and promotes early growth in *Arabidopsis thaliana*. *Proceedings of the National Academy of Sciences, USA* **99**, 17197–17202.

Igamberdiev AU, Hill RD. 2004. Nitrate, NO, and hemoglobin in plant adaptation to hypoxia: an alternative to classic fermentation pathways. *Journal of Experimental Botany* **55**, 2473–2482.

Maekawa T, Maekawa-Yoshikawa M, Takeda N, Imaizumi-Anraku H, Murooka Y, Hayashi M. 2009. Gibberellin controls the nodulation signaling pathway in *Lotus japonicus*. *The Plant Journal* **58**, 183–194.

Małolepszy A, Urbański DF, James EK, Sandal N, Isono E, Stougaard J, Andersen SU. 2015. The deubiquitinating enzyme AMSH1 is required for rhizobial infection and nodule organogenesis in *Lotus japonicus*. *The Plant Journal* **83**, 719–731.

Małolepszy A, Mun T, Sandal N, et al. 2016. The LORE1 insertion mutant resource. *The Plant Journal*. doi:10.1111/tpj.13243.

Mur LAJ, Santosa IE, Laarhoven LJJ, Holton NJ, Harren FJM, Smith AR. 2005. Laser photoacoustic detection allows in planta detection of nitric oxide in tobacco following challenge with avirulent and virulent *Pseudomonas syringae* pathovars. *Plant Physiology* **138**, 1247–1258.

Mur LAJ, Sivakumaran A, Mandon J, Cristescu S, Harren FJM, Hebelstrup KH. 2012. Haemoglobin modulates salicylate and jasmonate/ethylene-mediated resistance mechanisms against pathogens. *Journal of Experimental Botany* **63**, 4375–4387.

Nagata M, Murakami E, Shimoda Y, Shimoda-Sasakura F, Kucho K, Suzuki A, Abe M, Higashi S, Uchiumi T. 2008. Expression of a class 1 hemoglobin gene and production of nitric oxide in response to symbiotic and pathogenic bacteria in *Lotus japonicus*. *Molecular Plant-Microbe Interactions* **21**, 1175–1183.

Ott T, van Dongen JT, Günther C, Krusell L, Desbrosses G, Vigeolaas H, Bock V, Czechowski T, Geigenberger P, Udvardi MK. 2005. Symbiotic leghemoglobins are crucial for nitrogen fixation in legume root nodules but not for general plant growth and development. *Current Biology* **15**, 531–535.

Pajuelo E, Stougaard J. 2005. *Lotus japonicus* as a model system. In: Márquez A, ed. *Lotus japonicus* handbook. Dordrecht: Springer, 3–24.

Pauly N, Ferrari C, Andrio E, Marino D, Piardi S, Brouquisse R, Baudouin E, Puppo A. 2011. MtNOA1/RIF1 modulates *Medicago truncatula*–*Sinorhizobium meliloti* nodule development without affecting its nitric oxide content. *Journal of Experimental Botany* **62**, 939–948.

Perazzolli M, Dominici P, Romero-Puertas MC, Zago E, Zeier J, Sonoda M, Lamb C, Delledonne M. 2004. Arabidopsis nonsymbiotic hemoglobin AHb1 modulates nitric oxide bioactivity. *The Plant Cell* **16**, 2785–2794.



- Perry J, Brachmann A, Welham T, Binder A, Charpentier M, Groth M, Haage K, Markmann K, Wang TL, Parniske M.** 2009. TILLING in *Lotus japonicus* identified large allelic series for symbiosis genes and revealed a bias in functionally defective ethyl methanesulfonate alleles toward glycine replacements. *Plant Physiology* **151**, 1281–1291.
- Qu ZL, Zhong NQ, Wang HY, Chen AP, Jian GL, Xia GX.** 2006. Ectopic expression of the cotton non-symbiotic hemoglobin gene *GhHb1* triggers defense responses and increases disease tolerance in *Arabidopsis*. *Plant and Cell Physiology* **47**, 1058–1068.
- Sainz M, Pérez-Rontomé C, Ramos J, Mulet JM, James EK, Bhattacharjee U, Petrich JW, Becana M.** 2013. Plant hemoglobins may be maintained in functional form by reduced flavins in the nuclei, and confer differential tolerance to nitro-oxidative stress. *The Plant Journal* **76**, 875–887.
- Sasakura F, Uchiumi T, Shimoda Y, Suzuki A, Takenouchi K, Higashi S, Abe M.** 2006. A class 1 hemoglobin gene from *Alnus firma* functions in symbiotic and nonsymbiotic tissues to detoxify nitric oxide. *Molecular Plant-Microbe Interactions* **19**, 441–450.
- Shimoda Y, Nagata M, Suzuki A, Abe M, Sato S, Kato T, Tabata S, Higashi S, Uchiumi T.** 2005. Symbiotic rhizobium and nitric oxide induce gene expression of non-symbiotic hemoglobin in *Lotus japonicus*. *Plant and Cell Physiology* **46**, 99–107.
- Shimoda Y, Shimoda-Sasakura F, Kucho K, Kanamori N, Nagata M, Suzuki A, Abe M, Higashi S, Uchiumi T.** 2009. Overexpression of class 1 plant hemoglobin genes enhances symbiotic nitrogen fixation activity between *Mesorhizobium loti* and *Lotus japonicus*. *The Plant Journal* **57**, 254–263.
- Smagghe BJ, Hoy JA, Percifield R, et al.** 2009. Correlations between oxygen affinity and sequence classifications of plant hemoglobins. *Biopolymers* **91**, 1083–1096.
- Smagghe BJ, Trent III JT, Hargrove MS.** 2008. NO dioxygenase activity in hemoglobins is ubiquitous *in vitro*, but limited by reduction *in vivo*. *PLoS One* **3**, e2039.
- Tominaga A, Nagata M, Futsuki K, et al.** 2009. Enhanced nodulation and nitrogen fixation in the abscisic acid low-sensitive mutant *enhanced nitrogen fixation1* of *Lotus japonicus*. *Plant Physiology* **151**, 1965–1976.
- Trevaskis B, Watts RA, Andersson CR, Llewellyn DJ, Hargrove MS, Olson JS, Dennis ES, Peacock WJ.** 1997. Two hemoglobin genes in *Arabidopsis thaliana*: the evolutionary origins of leghemoglobins. *Proceedings of the National Academy of Sciences, USA* **94**, 12230–12234.
- Trinchant J-C, Rigaud J.** 1982. Nitrite and nitric oxide as inhibitors of nitrogenase from soybean bacteroids. *Applied and Environmental Microbiology* **44**, 1385–1388.
- Uchiumi T, Shimoda Y, Tsuruta T, et al.** 2002. Expression of symbiotic and nonsymbiotic globin genes responding to microsymbionts on *Lotus japonicus*. *Plant and Cell Physiology* **43**, 1351–1358.
- Urbański DF, Małolepszy A, Stougaard J, Andersen SU.** 2012. Genome-wide LORE1 retrotransposon mutagenesis and high-throughput insertion detection in *Lotus japonicus*. *The Plant Journal* **69**, 731–741.
- Vieweg MF, Hohnjec N, Küster H.** 2005. Two genes encoding different truncated hemoglobins are regulated during root nodule and arbuscular mycorrhiza symbioses of *Medicago truncatula*. *Planta* **220**, 757–766.
- Wally OSD, Mira MM, Hill RD, Stasolla C.** 2013. Hemoglobin regulation of plant embryogenesis and plant pathogen interaction. *Plant Signaling and Behavior* **4**, e25264.
- Watts RA, Hunt PW, Hvitved NA, Hargrove MS, Peacock WJ, Dennis ES.** 2001. A hemoglobin from plants homologous to truncated hemoglobins of microorganisms. *Proceedings of the National Academy of Sciences, USA* **98**, 10119–10124.
- Wopereis J, Pajuelo E, Dazzo FB, Jiang Q, Gresshoff PM, de Bruijn FJ, Stougaard J, Szczyglowski K.** 2000. Short root mutant of *Lotus japonicus* with a dramatically altered symbiotic phenotype. *The Plant Journal* **23**, 97–114.