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***Lotus japonicus* Contains Two Distinct ENOD40 Genes That Are Expressed in Symbiotic, Nonsymbiotic, and Embryonic Tissues**

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ENOD40, an early nodulin gene, has been postulated to play a significant role in legume root nodule ontogenesis. We have isolated two distinct ENOD40 genes from *Lotus japonicus*. The transcribed regions of the two ENOD40 genes share 65% homology, while the two promoters showed no significant homology. Both transcripts encode a putative dodecapeptide similar to that identified in other legumes forming determinate nodules. Both ENOD40 genes are coordinately expressed following inoculation of roots with *Mesorhizobium loti* or treatment with purified Nod factors. In the former case, mRNA accumulation could be detected up to 10 days following inoculation while in the latter case the accumulation was transient. High levels of both ENOD40 gene transcripts were found in nonsymbiotic tissues such as stems, fully developed flowers, green seed pods, and hypocotyls. A relatively lower level of both transcripts was observed in leaves, roots, and cotyledons. In situ hybridization studies revealed that, in mature nodules, transcripts of both ENOD40 genes accumulate in the nodule vascular system; additionally, in young seed pods strong signal is observed in the ovule, particularly in the phloem and epithelium, as well as in globular stage embryos.

Additional keywords: nodule organogenesis, symbiosis.

A group of soil bacteria that are collectively called rhizobia can cause the development of root nodules on their leguminous host plants. The developmental program of nodule initiation is triggered by signal molecules exchanged between the two symbionts, resulting in a complex temporal and spatial expression of several plant genes. Genes induced early during the symbiosis are defined as early noduleins (ENOD). Several lines of evidence suggest that early nodulin genes may play a

role in mediating rhizobial invasion and nodule organogenesis (Mylona et al. 1995). A number of early nodulin cDNA clones have been isolated and characterized, but the function of the encoded proteins still remains unknown. Among the early noduleins, ENOD40 is one of the most rapidly induced during the process of nodule organogenesis in both determinate and indeterminate nodule-forming legumes (Kouchi and Hata 1993; Yang et al. 1993; Crespi et al. 1994). The ENOD40 gene is first expressed, in response to rhizobial inoculation, in the root pericycle and subsequently in the dividing cortical cells. In mature, determinate root nodules, ENOD40 transcripts are detected in the pericycle of the vascular bundles (Kouchi and Hata 1993; Yang et al. 1993). A similar spatial pattern of expression is also observed during the ontogeny of indeterminate nodules (Asad et al. 1994; Crespi et al. 1994; Matvienko et al. 1995). Expression of ENOD40 has been reported in non-symbiotic tissues, such as stems (Kouchi and Hata 1993; Yang et al. 1993; Asad et al. 1994), lateral and adventitious root primordia (Papadopoulou et al. 1996) and leaf and stipule primordia (Frang and Hirsch 1998; Corich et al. 1998). ENOD40 gene homologues have also been identified in the nonlegumes tobacco (Matvienko et al. 1996) and rice (Kouchi et al. 1999). Expression analysis studies revealed that ENOD40 transcripts were detected only in the early developmental stages of stem vascular bundles of rice (Kouchi et al. 1999).

In a number of legumes, transcription of the ENOD40 gene could be induced in root tissues in response to purified Nod factors (Minami et al. 1996), cytokinin (van Rhijn et al. 1997), and chitin pentamers (Minami et al. 1996), and during arbuscular mycorrhizae infection (van Rhijn et al. 1997). ENOD40 transcripts were also found in spontaneous nodules (Crespi et al. 1994) and in empty nodules induced by mutant bacteria (Yang et al. 1993; Asad et al. 1994; Crespi et al. 1994).

Results obtained with transgenic *Medicago sativa* suggest that ENOD40 may be involved in the initiation of nodule formation by triggering cortical cell divisions (Charon et al. 1997). *M. sativa* embryos over-expressing ENOD40 developed into teratomas, whereas expression of an antisense ENOD40 construct in *M. sativa* explants resulted in arrest of callus growth and loss of embryo formation (Crespi et al. 1994). Moreover, over-expression of ENOD40 in transgenic

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Nucleotide sequences have been submitted to the EMBL database as accession nos. AJ271787 (*Lj*ENOD40-1) and AJ271788 (*Lj*ENOD40-2).

Medicago truncatula plants accelerates nodulation upon inoculation with *Sinorhizobium meliloti* (Charon et al. 1999).

The ENOD40 transcripts are characterized by the absence of a long open reading frame (ORF), which led to the hypothesis that ENOD40 genes code for an untranslated RNA. However, comparison of the potential ORFs of the ENOD40 gene transcripts identified so far revealed the presence of a small ORF coding for a highly conserved, small peptide 10 to 13 amino acids long (Vijn et al. 1995b; Kouchi et al. 1999).

In this study, we demonstrate that there are two distinct ENOD40 genes present in *Lotus japonicus* that are expressed in both symbiotic and nonsymbiotic tissues. In situ hybridization studies show that ENOD40 transcripts are localized in the vascular system of nodular and ovular tissues as well as in the globular stage embryos.

RESULTS

L. japonicus contains two distinct ENOD40 genes.

Several cDNA clones were isolated from a cDNA library prepared from 21-day-old *L. japonicus* nodules, with a cloned polymerase chain reaction (PCR) fragment (designated as *LjENOD40p*) of *L. japonicus* genomic DNA as a probe. The nucleotide sequence of the identified cDNA clones revealed that they all share identical sequences and showed high homology to the soybean ENOD40 genes. The largest cDNA, of 550 bp, excluding the polyA tail, was designated as *LjENOD40-1c*. To identify the total number of ENOD40 genes, genomic DNA from *L. japonicus* was digested with *EcoRI* and/or *HindIII* and subjected to Southern blot analysis. The data obtained revealed the presence of two *EcoRI* hybridizing bands of about 2.4 and 3.2 kb, respectively (data not shown).

To isolate and characterize the two putative ENOD40 genes, several genomic clones were isolated by screening an *L. japonicus* genomic library. Southern blot analysis of genomic *L. japonicus* DNA and representative genomic lambda clones indicated that the latter contained either of both hybridizing *EcoRI* fragments from the plant genome (data not shown). Nucleotide sequence analysis of both *EcoRI* fragments obtained from the lambda phages (Fig. 1A) revealed that the 2.4-kb *EcoRI* fragment contained sequences identical to the *LjENOD40-1c* cDNA. In contrast, the genomic 3.2-kb *EcoRI* fragment (Fig. 1A) contained sequences that are homologous but not identical to *LjENOD40-1c*. Regions of the second ENOD40 gene (designated as *LjENOD40-2*) that exhibited low homology with the respective regions of *LjENOD40-1c* were amplified, cloned, named *LjENOD40-2S*, and used as a probe to screen a cDNA library prepared from 9-day-old *L. japonicus* roots inoculated with *Mesorhizobium loti*. Several cDNA clones were identified and their nucleotide sequence was determined. Aside from variation in length, the nucleotide sequence of all the isolated cDNA clones was identical to that of the *LjENOD40-2* clone. The cDNA with the largest insert was designated as *LjENOD40-2c*. The transcription start of both ENOD40 genes was determined by 5'-rapid amplification of cDNA ends (5' RACE). Two distinct 5' ends were identified for *LjENOD40-1* transcripts while only one was identified for *LjENOD40-2* (Fig. 1B). Comparison of the nucleotide sequences of the two *L. japonicus* ENOD40 genes demonstrated that they share only 65% identity (Fig.

1B). It can be noted that a 55-bp insertion is present in the *LjENOD40-2* gene. A 75-bp insertion has also been detected in the alfalfa ENOD40 gene (Crespi et al. 1994). Sequence comparison revealed that the proximal sequences of the two promoter regions showed no significant sequence homology (Fig. 1B).

The percentage of nucleotide sequence similarity of the two *L. japonicus* ENOD40 genes, compared with those reported in other legumes, indicates that, within the legume family, two major ENOD40 groups have evolved, which can be designated as group I and group II (Fig. 2A). The legumes forming determinate nodules are clustered in group I and those forming indeterminate ones in group II. Interestingly, within group I, the two *L. japonicus* ENOD40 genes belong to different clades.

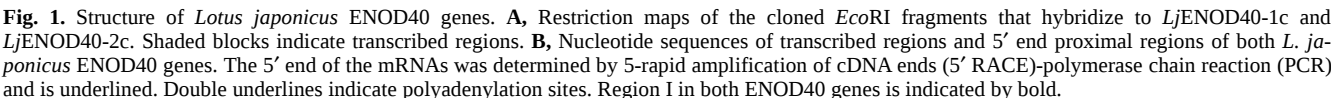
It has been demonstrated that all known ENOD40 genes in legumes contain two conserved regions, named regions I and II. Region I contains a small ORF encoding a peptide of 12 or 13 amino acids in determinate or indeterminate nodule forming legumes, respectively (Vijn et al. 1995b). Analysis of the nucleotide sequence of the two *L. japonicus* cDNA clones demonstrated that both contained the proposed regions I and II. In both ENOD40 gene transcripts, region I encodes for the proposed conserved 12 amino acids peptide (Figs. 1B and 2B). It should be noted that the pI values of the predicted peptides differ significantly. Interestingly, the pI value of the peptide encoded by the *LjENOD40-1c* clone appears to be similar to that of *Sesbania rostrata* and common bean, whereas the pI value of the peptide encoded by the *LjENOD40-2c* clone is similar to that of pea and alfalfa (Fig. 2B).

Both ENOD40 genes are induced by *M. loti* and purified LCOs.

The accumulation of both ENOD40 genes transcripts was examined after inoculation with *M. loti* or treatment with purified lipo-chitin oligosaccharides (LCOs). Accumulation of ENOD40 transcripts was examined up to 10 days post inoculation with *M. loti* and the two ENOD40 transcripts showed the same dynamics of expression (Fig. 3A). Treatment with purified *M. loti* LCOs induced, at 5 days post treatment, similar dynamics of accumulation of both ENOD40 genes, at levels comparable to those observed 5 days after inoculation with bacteria. However, at 10 days post treatment, the accumulation of transcripts was markedly reduced to basal levels (Fig. 3A). It should also be pointed out that no visible, nodule-like structures were observed 10 days after the treatment of the young seedlings with LCOs, in contrast with what was previously reported in the case of *Lotus preslii* (Lopez-Lara et al. 1995).

Expression of ENOD40 genes in nonsymbiotic tissues of *L. japonicus*.

Several groups have reported that in legumes the ENOD40 gene, in addition to its expression in symbiotic organs, is also expressed in nonsymbiotic organs (Crespi et al. 1994; Asad et al. 1994; Papadopoulou et al. 1996; Corich et al. 1998). Thus, it was of interest to investigate whether both ENOD40 genes are also expressed in various plant organs. To test this, a semiquantitative reverse transcription (RT)-PCR analysis was employed. The results indicated that the mRNAs isolated from mature nodules, hypocotyls, and young stems contained the highest levels of both ENOD40 gene transcripts (Fig. 3B).



Furthermore, high levels of both transcripts were present in mRNA prepared from fully developed flowers and green seed pods (Fig. 3B). In contrast, relatively lower levels of both transcripts were detected in roots and germinated cotyledons, whereas in leaves the expression of both ENOD40 genes was hardly detectable (Fig. 3B).

Spatial localization of ENOD40 gene transcripts.

The spatial localization of ENOD40 gene transcripts was investigated by an in situ hybridization approach. At 10 days after inoculation with *M. loti*, *LjENOD40-1* is highly expressed in the developing vascular bundles (Fig. 4A and B), as was previously reported (Kouchi and Hata 1993; Yang et al.

1993). Interestingly, *LjENOD40-2* gene transcripts showed a similar expression pattern in 10-day-old nodules (Fig. 4C), when clone *LjENOD40-2S* was used to transcribe the antisense RNA probe. In 21-day-old nodules, *LjENOD40-1* gene transcripts accumulate in the vascular tissue surrounding the nodule (Fig. 4D and E). Additionally, in green young pods, *LjENOD40-1* is up-regulated in the ovules and very strong hybridization signal is observed in the phloem tissue (Fig. 4F and G) and epithelium cells (Fig. 4F). High expression was also observed at the globular stage embryo (Fig. 4H). In the seed pod tissues surrounding the ovule, no hybridization signal above background was detected. As a negative control, sections were hybridized to sense RNA probe transcribed from *LjENOD40-1c* clone (Fig. 4J–M). In this case, no significant hybridization signal was detected.

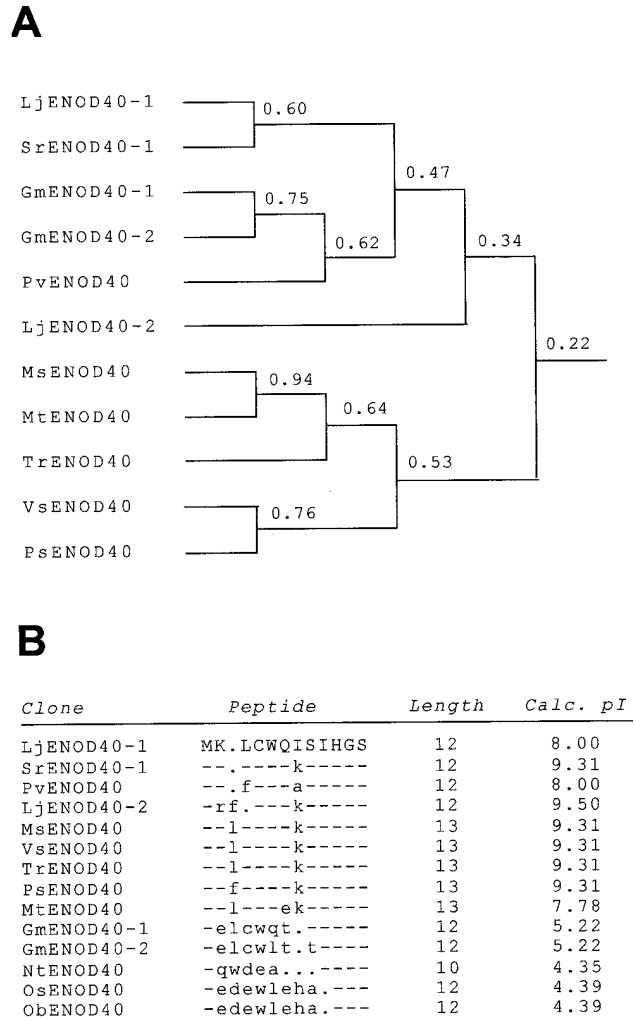


Fig. 2. Phylogenetic relationship of ENOD40 genes from various legumes and comparison of putative encoded oligopeptides. **A**, Phylogenetic tree based on ENOD40 cDNA sequences from various leguminous plants. **B**, Comparison of amino acid sequences encoded by region I of ENOD40 genes from various plants, together with predicted pI values. Plant species and DDBJ, EMBL, and GenBank data base accession nos.: *Lj*, *Lotus japonicus* (this work); *Gm*, *Glycine max* (a, D13503; b, D13504); *Sr*, *Sesbania rostrata* (Y12714); *Pv*, *Phaseolus vulgaris* (X86441); *Ms*, *Medicago sativa* (X80263); *Vs*, *Vicia sativa* (X83683); *Tr*, *Trifolium repens* (AJ000268); *Ps*, *Pisum sativum* (X81064); *Mt*, *Medicago truncatula* (X80264); *Nt*, *Nicotiana tabacum* (X98716); *Os*, *Oryza sativa* (ABO24054); *Ob*, *Oryza brachyantha* (ABO24055).

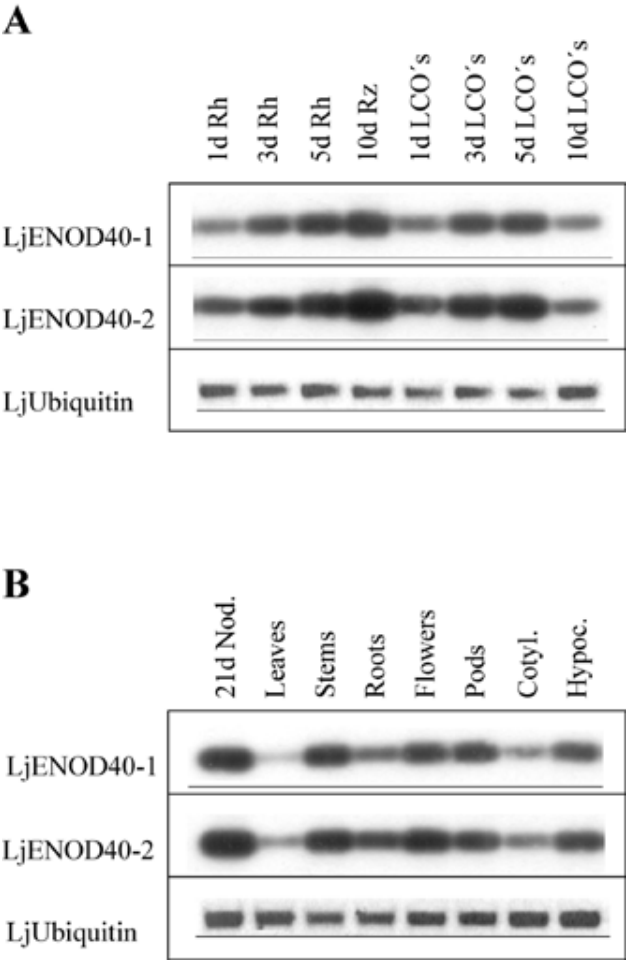


Fig. 3. Accumulation of *LjENOD40* genes transcripts in symbiotic and nonsymbiotic tissues of *Lotus japonicus* and after treatment with *Mesorhizobium loti* purified lipo-chitin oligosaccharides (LCOs). Total RNAs from the various tissues were subjected to reverse transcription-polymerase chain reaction (RT-PCR) analysis with *LjUbiquitin* as internal control. Amplification products were analyzed on 1.5% agarose gels, blotted to nylon membranes, and probed with *LjENOD40-1c* and *LjENOD40-2c* respectively. **A**, Lanes 1–4, *L. japonicus* (Gifu B-129) roots 1, 3, 5, and 10 days after inoculation with *M. loti* (strain E1R.pMP2112). Lanes 5–8, *L. japonicus* (Gifu B-129) roots 1, 3, 5, and 10 days after treatment with purified *M. loti* LCOs. **B**, Lanes 1–8, 21-day-old *L. japonicus* nodules, young leaves, young stems, roots, flowers, green seed pods, germinated cotyledons, and hypocotyls, respectively.

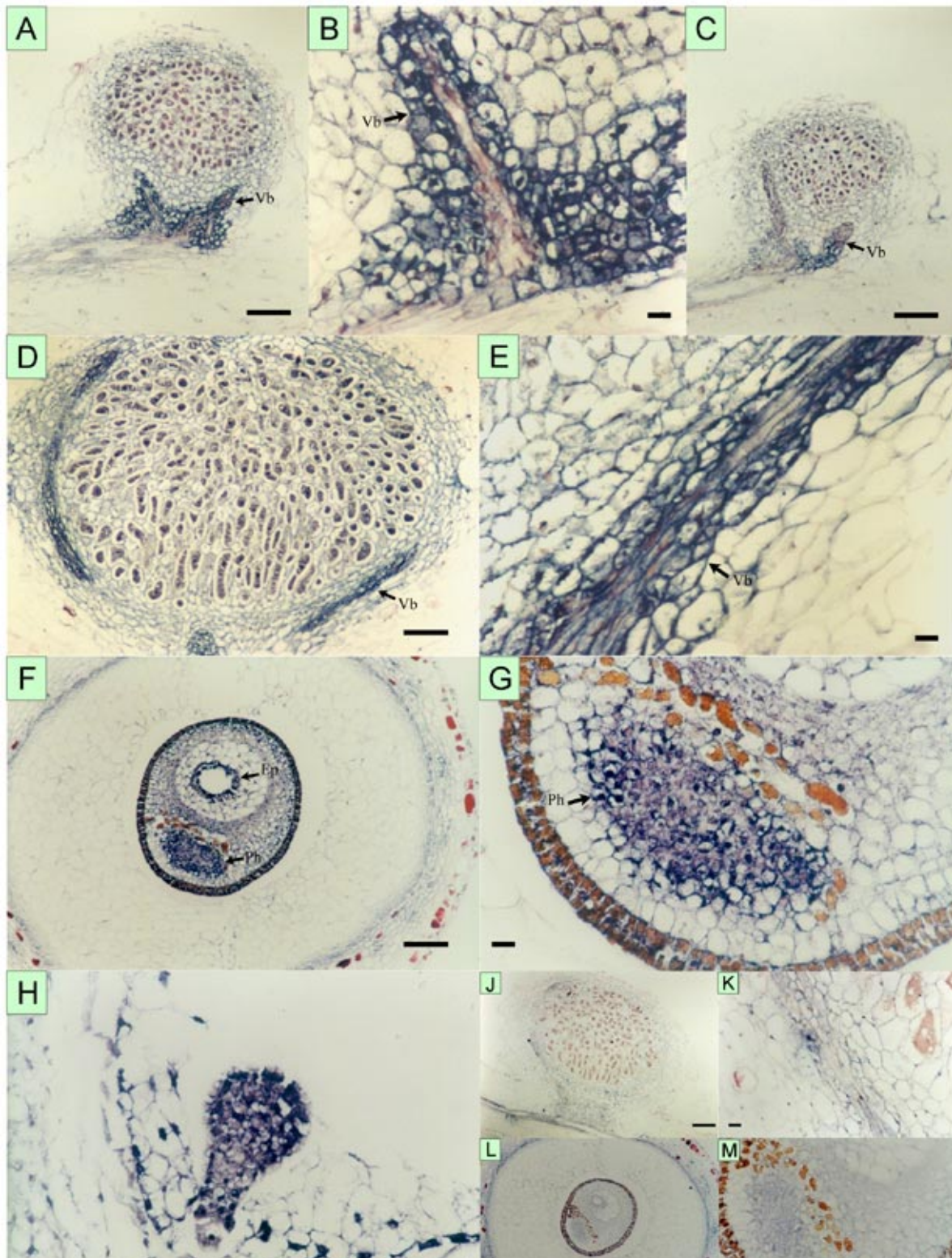


Fig. 4. In situ localization of *LjENOD40* genes transcripts in *Lotus japonicus* (Gifu B-129) root nodules, young seed pods, and embryos. Hybridization signal is visible as blue-purple precipitate. **A** and **B**, *LjENOD40-1* transcripts in 10-day-old nodules are present in the developing vascular bundles (Vb). **C**, In nodules of the same age *LjENOD40-2* is also expressed in developing vascular tissue. **D** and **E**, In mature 21-day-old nodules *LjENOD40-1* is highly expressed in vascular tissues of nodule. **F** and **G**, In young seed pods *LjENOD40-1* transcripts are clearly detected in ovule. Elevated level of expression is observed in phloem tissue (Ph) of ovule and epithelium cells (Ep). No visible hybridization signal is detected in pod tissues surrounding ovule. **H**, Markedly high expression level of *LjENOD40-1* is detected in globular stage embryos. As a negative control, sections of (**J** and **K**) 21-day-old nodules and (**L** and **M**) young seed pods were hybridized to sense RNA transcribed from *LjENOD40-1c* clone. In this case, no significant hybridization signal is visible. Bars = (**A,C,D,F,J,L**) 100 μ m or (**B,E,G,H,K,M**) 10 μ m.

DISCUSSION

The fact that genomic clones and transcripts for the two distinct ENOD40 genes have been found in the diploid *L. japonicus*, provides strong evidence that there are at least two functional ENOD40 genes. The presence of two ENOD40 genes has also been proposed for the hexaploid soybean (Minami et al. 1996), the tetraploid alfalfa (Frang and Hirsch 1998), and *S. rostrata* (Corich et al. 1998). However, in the case of alfalfa, the two genes including a large part of the promoter region are identical, while the two soybean ENOD40 genes share a 94% homology, thus not excluding the possibility that both cases may represent two variants of the same allele. The SrENOD40-1 is 84.5% identical to SrENOD40-2. However, SrENOD40-2 appears to lack the sequences coding for the dodecapeptide (Corich et al. 1998). Since the homology of the two *L. japonicus* ENOD40 cDNA clones is only 65% it could be argued that a second, distinct ENOD40 gene in other legumes has not yet been found, possibly due to low homology with the identified one.

Molecular, phylogenetic analysis inferred from sequence data of the available ENOD40 genes revealed that ENOD40 genes form two groups among the legumes. An almost identical grouping of legumes was obtained by Doyle et al. (1998), with *rbcL* sequences. Within group I, the two *L. japonicus* ENOD40 genes belong to different clades; the *Lj*ENOD40-1 gene showed homology of about 78 and 72% to the soybean ENOD40-1 and common bean ENOD40 genes, respectively, and only about 65% to the *Lj*ENOD40-2 gene (Fig. 2A). The sequence divergence of the two *L. japonicus* ENOD40 genes can be partly attributed to insertion sequences, a feature also found in ENOD40 cDNA clones isolated from legumes forming indeterminate nodules (Crespi et al. 1994). These data also suggest that ENOD40 gene duplication events have taken place early in legume evolution.

Both *L. japonicus* ENOD40 transcripts have the capacity to code the putative 12 amino acid peptide found in legumes that form determinate nodules. Comparison of the putative peptides encoded by ENOD40 genes in different legume and nonlegume plants indicated a diverse calculated pI value (Fig. 2B). This observation raises the question of whether these peptides may require post-translational modification to be functional and/or that formation of functional complexes with other compounds is necessary for the peptides to be functional in vivo. The sort mating factor peptide, such as the decapeptide from *Rhodospiridium toruloids*, has been shown to require farnesylation of the C-terminal amino acids to be functional (Rodriguez et al. 1999). Despite the fact that region II is highly conserved at the nucleotide level between the two *L. japonicus* ENOD40 genes, it lacks a conserved typical ORF. This is also the case for all the ENOD40 homologues studied so far (Kouchi et al. 1999), indicating that the translation of this region is not necessary for it to be biological active (Charon et al. 1997).

Expression analysis, with in situ hybridization and semi-quantitative RT-PCR, showed that the two *L. japonicus* ENOD40 genes are co-expressed in the same developmental stage and in the same cell types during nodule organogenesis. Since both genes are induced by purified LCOs, it is plausible to assume that similar signal transduction pathways triggered by the Nod factors might be involved in the activation of

equivalent regulatory elements present in the promoters of both genes. The regulatory elements that are required for the induction of both pea ENOD12 (Vijn et al. 1995a) and ENOD40 (Fang and Hirsch 1998) by Nod factors appear to be located within 200 and 600 bp, respectively, upstream of the translation start. If identical factors can act on both *L. japonicus* ENOD40 genes, one would expect to find sequence motifs in both genes that are similar. Alternatively, the coordinated induction of the two ENOD40 genes may be mediated by two types of Nod receptors similar to those suggested for alfalfa (Ardourel et al. 1994).

The expression patterns of ENOD40 genes in *L. japonicus* root nodules followed basically the ones described for soybean (Kouchi and Hata 1993; Yang et al. 1993) and common bean (Papadopolou et al. 1996). In fully developed nodules, ENOD40 is detected in specific cells of the pericycle of the nodule vascular bundles. Similarly, in *L. japonicus* embryonic tissues, high levels of expression of ENOD40-1 gene are also observed in the ovules and specifically the strongest signal was observed in the ovule phloem and ovule epithelium cells, as well as in the embryo itself. In rice, expression of the *Os*ENOD40 gene is coupled to specific developmental stages of the stem's lateral vascular bundles (Kouchi et al. 1999). These data, taken together, are in accordance with the hypothesis that ENOD40 gene products mediate the transport of specific metabolites or factors necessary for cell development and growth, as has been proposed before (Papadopolou et al. 1996). The possible involvement of ENOD40 gene products in organogenetic processes in nonsymbiotic tissues was shown by antisense RNA experiments in *M. sativa* explants that resulted in arrest of callus growth and loss of embryo formation while, in plants over-expressing ENOD40, embryos developed into teratomas (Crespi et al. 1994). The expression of ENOD40 in developing embryos, as well as in ovule tissues, of *L. japonicus* adds further proof to the suggestion that the ENOD40 gene is a plant gene involved in organogenesis (Crespi et al. 1994; Asad et al. 1994; Papadopolou et al. 1996).

The presence of both ENOD40 gene transcripts in nonsymbiotic tissues of legumes and the occurrence of homologues of ENOD40 in nonleguminous plants such as tobacco (Matvienko et al. 1996) and rice (Kouchi et al. 1999), could mean that the regulatory elements of these genes may be directly activated by factors that are recognizing endogenous LCO-like compounds or other signal molecules that functionally resemble LCOs. This hypothesis is suggested by the biological effects of LCOs on nonlegumes such as carrots (De Jong et al. 1993) or rice (Reddy et al. 1998). It is also possible that the ENOD40 genes are indirectly triggered by Nod factors via an effect on the levels of common plant hormones such as auxin. Evidence for this is provided by the observation that microtargeting of Nod factors or O-acetylated chitin oligomers into the root cortex appears to elicit changes in the characteristics of the root auxin balance during nodule initiation in clover (Mathesius et al. 1998). The information obtained in this study will be very important to further address these hypotheses about the regulation of the ENOD40 genes in the *L. japonicus* system. Analysis of transgenic *L. japonicus* plants containing a reporter gene whose expression is driven by these promoters would allow further insight into the mechanism of ENOD40 gene induction. For this purpose, constructs of both

LjENOD40 promoters fused to a *gus-gfp* gene (Quaedvlieg et al. 1998) have been introduced into *L. japonicus* with the *Agrobacterium tumefaciens* system and await further analysis after transgenic seeds have been obtained.

MATERIALS AND METHODS

Plant material and growth conditions.

L. japonicus (Gifu B-129) seeds were kindly provided by Jens Stougaard (University of Aarhus, Denmark). All plants were grown in a controlled environment with a 18 h/ 6h day/night cycle, a 22/18°C day/night regime and 70% humidity (Handberg and Stougaard 1992). Prior to germination, seeds were scarified by treatment with H₂SO₄ for 5 min and sterilized for 20 min with 2% NaOCl–0.02% Tween 20. Seeds were pre-germinated at 18°C in the dark for 72 h and the small plants were grown in Hoagland nutrient solution. For the inoculation with *Rhizobium* spp., 72-h seedlings were inoculated with a 0.1 OD₆₀₀ suspension of the *M. loti* strain E1R.pMP2112 and the plants were grown in nitrogen-free B&D nutrient solution (Broughton and Dilworth 1971). Nod factors were isolated from the *M. loti* strain E1R.pMP2112 by published methods (Lopez-Lara et al. 1995), while the method of application and the concentration of the Nod factors were the same as published for *L. preslii* (Lopez-Lara et al. 1995).

Isolation of *LjENOD40* cDNA and genomic clones.

For the amplification of *ENOD40* homologous sequences from *L. japonicus*, two degenerate primers designated 40D1F [5'-ATCCATGGTTCTT(AG)AAGAAG-3'] and 40D2R [5'-(GC)C(TC)TTTT(TC)GTGACTTGCCG-3'] were designed with previously published *ENOD40* sequences from various leguminous plants. The PCR product was cloned in pBlueScript KS⁺ plasmid vector (Stratagene, La Jolla, CA), resulting in a plasmid named p*LjENOD40p*. The nucleotide sequence of the insert was determined by the dideoxy chain termination method (Sanger et al. 1997).

A λ ZAPII *L. japonicus* polyT-primed cDNA library from 21-day-old nodules and a λ fixII *L. japonicus* genomic library was screened by plaque hybridization at 65°C, with the ³²P-dCTP-labeled insert of the *LjENOD40p* clone (both libraries were kindly provided by Jens Stougaard). Phage purification and in vitro excision of pBlueScript SK⁺ plasmids from the λ ZAPII library were performed according to standard methods (Stratagene) and the nucleotide sequence was determined as above. The purified phages from the λ FIXII genomic library, giving positive hybridization signal were digested with various restriction enzymes blotted and hybridized with the ³²P-dCTP-labeled insert of the *LjENOD40p* clone used as a probe. The hybridizing *Eco*RI fragments were subcloned in pBlueScript KS⁺ plasmid vector and the nucleotide sequence was determined.

For the isolation of a cDNA clone of the *LgENOD40-2* gene, a specific region from the genomic clone (showing 35% homology with *LgENOD40-1* clone), was amplified by PCR with the oligonucleotides *Lg40-2F* (5'-GAATGTGTGA GCGGGTCAGG-3') and *Lg40-2R* (5'-ACTTCTTCTACTC ATAAAGA-3'). The amplification product was cloned in pBlueScript KS⁺ and the clone was designated as p*LjENOD40-2S*. The insert of this clone was labeled with ³²P-dCTP and used as a probe to screen an *L. japonicus* random-

primed λ ZAPII cDNA library from 9-day-old roots inoculated with *M. loti* (kindly provided by Jens Stougaard). Phage purification and in vitro excision of the pBlueScript SK⁺ plasmids from the λ ZAPII library were performed according to standard methods (Stratagene) and the nucleotide sequence was determined.

Detection of *LjENOD40-1* and *LjENOD40-2* transcripts with an RT-PCR assay.

All the *L. japonicus* (Gifu B-129) tissues of interest and young roots inoculated with *M. loti* (strain E1R.pMP2112) or treated with purified *M. loti* Nod factors were harvested and ground in liquid nitrogen. Total RNA was isolated according to Brusslan and Tobin (1992) and quantified by spectrophotometry and agarose gel electrophoresis. Prior to RT-PCR, the total RNA samples were treated with DNase I (Promega, Madison, WI) at 37°C for 10 min for the elimination of contaminating genomic DNA. The Titan One Tube RT-PCR System (Boehringer Mannheim, Mannheim, Germany) was used for the reverse transcription and amplification of *ENOD40* transcripts. Three primers were designed: one common primer for both transcripts, named *Lj40-C* (5'-CCTACTCATCT GCAGAACTG-3') and two specific primers designated as *Lj40-1F* (5'-GGAGGTATGCTCAAACAT TC-3') and *Lj40-2F* (5'-CAAACTCGTTATGTTGCGG-3'). One hundred nanograms of total RNA treated with DNase I was used for each reaction and the different RNA preparations were normalized by the parallel amplification of the constitutively expressed gene *LjUbiquitin* with *LjUBQ-F* (5'-ATGCAGATCTTTTGT GAAGAC-3') and *LjUBQ-R* (5'-ACCACCACGGAAGA CGGAG-3') primers. Under our experimental conditions an exponential increase of the *ENOD40* RT-PCR products was still observed after 25 cycles (94°C/1 min, 52°C/1 min, 68°C/1 min), so we performed all the reactions under these conditions to obtain semiquantitative results. The RT-PCR products were analyzed by 1.5% agarose gel electrophoresis, blotted, and hybridized to *LjENOD40-1* and *LjENOD40-2* cDNA probes.

Determination of transcription start with a RACE-PCR assay.

Total RNA was isolated from 21-day-old nodules as described above and a 5' RACE-PCR was performed with the SMART 5' RACE kit from Clontech (Westburg, NL). Two gene-specific oligonucleotides were used as primers, one for each *ENOD40* transcript. *Lj40-1Rc* (5'-GAATGTTTGAGCA TACCTCCATGGTG-3') was used for the amplification of *LjENOD40-1* specific 5'-end sequences and *Lj40-2Rc* (5'-GGTATCCCGGTGATTACTAACTCGG-3') was used in the case of *LjENOD40-2*. The RACE-PCR products were cloned in pBlueScript KS⁺ plasmid vector (Stratagene) and the nucleotide sequence was determined.

In situ hybridization.

L. japonicus nodules harvested at 10 and 21 days after inoculation of roots with *M. loti* (strain E1R.pMP2112) and young pods were fixed in 4% paraformaldehyde supplemented with 0.25% glutaraldehyde in 10 mM sodium phosphate buffer pH 7.4 for 4 h in a vacuum aspirator (Papadopolou et al. 1996). Fixed tissues were dehydrated through ethanol series, embedded in paraffin, and 7- μ m-thin sections were cut as

described by Yang et al. (1993). Antisense and sense RNA probes labeled with digoxigenin (DIG)-11-rUTP (Boehringer Mannheim) were transcribed from the *LjENOD40-1* and *LjENOD40-2S* clones with the T3 and T7 promoters of pBlueScript SK⁺. The RNA probes were hybridized to the sections and hybridization signals were visualized with anti-DIG antibodies conjugated with alkaline phosphatase (Kouchi and Hata 1993).

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