Large-Scale Analysis of Putative Soybean Regulatory Gene Expression Identifies a Myb Gene Involved in Soybean Nodule Development\textsuperscript{1[W][OA]}

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Nodulation is the result of a symbiosis between legumes and rhizobial bacteria in soil. This symbiosis is mutually beneficial, with the bacteria providing a source of nitrogen to the host while the plant supplies carbon to the symbiont. Nodule development is a complex process that is tightly regulated in the host plant cell through networks of gene expression. In order to examine this regulation in detail, a library of quantitative reverse transcription-polymerase chain reaction primer sets was developed for a large number of soybean (Glycine max) putative regulatory genes available in the current expressed sequence tag collection. This library contained primers specific to soybean transcription factor genes as well as genes involved in chromatin modification and translational regulation. Using this library, we analyzed the expression of this gene set during nodule development. A large number of genes were found to be differentially expressed, especially at the later stages of nodule development when active nitrogen fixation was occurring. Expression of these putative regulatory genes was also analyzed in response to the addition of nitrate as a nitrogen source. This comparative analysis identified genes that may be specifically involved in nitrogen assimilation, metabolism, and the maintenance of active nodules. To address this possibility, the expression of one such candidate was studied in more detail by expressing in soybean roots promoter β-glucuronidase and green fluorescent protein fusions. This gene, named Control of Nodule Development (CND), encoded a Myb transcription factor gene. When the CND gene was silenced, nodulation was reduced. These results, associated with a strong expression of the CND gene in the vascular tissues, suggest a role for CND in controlling soybean nodulation.

Nodulation involves the intimate relationship between soil bacteria (rhizobia) and legume plants, which results in the formation of a novel organ, the nodule, in which the bacteria reside and provide a steady source of nitrogen to the plant. Control of the initial host-symbiont interaction and subsequent nodule development is complex and appears to require a continuous exchange of chemical signals between the partners. For example, the compatible symbiont initially recognizes flavonoid molecules released by the host. This, in turn, leads to the production of a specific lipochitin nodulation signal (Nod factor) that is required for rhizobial infection and also triggers the development of the nodule primordium (Oldroyd and Downie, 2008).

Primarily through an examination of plant mutants defective in nodulation, the basic steps in the Nod factor signaling pathway have been elucidated (Oldroyd and Downie, 2008). Initial Nod factor recognition is mediated through two LysM receptors that likely directly interact with the compatible Nod factor (Amor et al., 2003; Madsen et al., 2003; Radutoiu et al., 2003). The resulting signaling cascade ultimately leads to activation of specific transcription factors (TFs), a few of which have been identified. For example, the Medicago truncatula MtNSP1 and MtNSP2 genes encode two GRAS family TFs (Catoira et al., 2000; Oldroyd and Long, 2003; Kalo et al., 2005; Smit et al., 2005) that are essential to root hair infection. MtERN, a member of the AP2-EREBP family (Middleton et al., 2007), was shown to play a key role in the initiation and maintenance of rhizobial infection. In Lotus japonicus, the NIN gene encodes a putative TF gene (Schauser et al., 1999). Mutants in the L. japonicus NIN gene, or the Pisum sativum ortholog Sym35, fail to support rhizobial infection and do not show cortical cell division upon inoculation (Schauser et al., 1999; Borisov et al., 2003). In contrast, the L. japonicus astray mutant exhibits hypernodulation. The ASTRAY gene encodes a bZIP TF (Nishimura et al., 2002).

The expression of some of these key TF genes is affected by rhizobial inoculation. For example, NSP2,
ERN, and NIN expression was induced in response to root inoculation by the compatible symbiont (Kalo et al., 2005; Marsh et al., 2007; Middleton et al., 2007). Such results suggest that TFs involved in nodulation may be identified by directly analyzing their transcriptional response to inoculation. Based on such facts, large-scale DNA microarray studies identified a variety of putative TF genes that were differentially expressed during nodule initiation and development (El Yahyaoui et al., 2004; Kouchi et al., 2004; Benedito et al., 2008; Brechenmacher et al., 2008). For example, in the study of El Yahyaoui et al. (2004), 24 M. truncatula TF genes were found to respond to rhizobial inoculation. Among these genes was the M. truncatula HAP2-1 gene, a member of the CCAAT-binding TF family. Combier et al. (2006) showed that MtHAP2-1 expression was regulated through the action of a specific microRNA (miR169). Boualem et al. (2008) identified a second microRNA in M. truncatula, miR166, targeting at least three class III HD-ZIP TF genes (MtCN1, MtCN2a, and MtHB8). The down-regulation of these three genes affected the number of infection initiation events in the symbiosis between M. truncatula and Sinorhizobium meliloti. Collectively, these studies reflect the important roles of TF genes in nodulation.

Due to the low global expression level of many TF genes, DNA microarrays lack the sensitivity to accurately measure their expression (Czechowski et al., 2004; Libault et al., 2007). Czechowski et al. (2004) created a resource that utilized highly sensitive quantitative reverse transcription (qRT)-PCR to accurately analyze the expression of more than 2,000 Arabidopsis (Arabidopsis thaliana) TF genes. This study clearly demonstrated that the use of such qRT-PCR primer libraries provides 100- to 1,000-fold greater sensitivity in accurately measuring TF gene expression compared with DNA microarray technologies. Since this initial paper, other primer libraries have been developed for the various putative regulatory genes in Oryza sativa (Caldana et al., 2007) and Medicago truncatula (Kakar et al., 2008). These libraries, as well as the original Arabidopsis library, were used in large-scale analyses to identify putative regulatory genes that are differentially expressed in different plant tissues, in response to pathogens and microbial elicitors, and after treatment with nitrogen and phosphorous (Czechowski et al., 2004; Scheible et al., 2004; McGrath et al., 2005; Caldana et al., 2007; Libault et al., 2007; Morcuende et al., 2007; Kakar et al., 2008; Verdier et al., 2008; Gruber et al., 2009).

In this study, by mining public EST databases, 2,586 putative soybean (Glycine max) regulatory genes (i.e. TF genes as well as those involved in chromatin modification and translational regulation) were identified. PCR primers were designed and validated to create 1,149 specific primer sets, which can be used to specifically quantify gene expression by qRT-PCR. To demonstrate the usefulness of this resource, the expression of these genes was profiled during soybean nodule development by comparing nodulated and nodule-free roots. The results showed that 126 putative regulatory genes were differentially regulated during nodulation, including putative orthologs of MtHAP2.1 and MtERN. Most of these genes were regulated at the later stages when active nitrogen fixation is occurring. Further analysis distinguished those putative regulatory genes strictly regulated during nodulation, after nitrogen starvation, and/or those responsive to the addition of nitrate as a nitrogen source. Control of Nodule Development (CND), a Myb TF gene, was studied in more detail through the construction of GUS and GFP promoter fusions. Silencing of CND resulted in a decrease in nodule numbers. Our data clearly indicate that a specific, temporal pattern of putative regulatory gene expression correlates with nodule development.

RESULTS AND DISCUSSION
Identification of 2,586 Putative Soybean Regulatory Genes in Public EST Databases
To begin compiling a library of putative regulatory genes in soybean, the sequences of predicted TF genes in Arabidopsis (2,290 and 2,250 predicted TF transcripts from the Database of Arabidopsis Transcription Factors [http://datf.cbi.pku.edu.cn/] and the Plant Transcription Factor Database [http://plntfdb.bio.uni-potsdam.de/v2.0/index.php?sp_id=ATH], respectively) were used for similarity searches to identify homologous soybean genes. Because the soybean genome sequences were not available at the time of the design of the primer sets, the identified sequences were compared against the National Center for Biotechnology Information G. max Unigene database (with BLAST e-value ≤ 1e-12) to identify homologous soybean sequences. In addition, The Institute for Genome Research soybean Gene Indices (http://www.tigr.org/tigr-scripts/tgi/T_index.cgi?species=soybean) and soybean EST databases (http://www.ncbi.nlm.nih.gov/dbEST) were mined to identify 1,321 tentative consensus sequences and 1,043 ESTs, respectively, that are annotated as regulatory genes. Finally, the annotation of the soybean Affymetrix arrays (Robert Goldberg, personal communication) was used to identify additional putative soybean regulatory genes. Altogether, after identifying and removing redundant sequences between the different databases, these different approaches identified 2,586 transcripts (Supplemental Table S1). In cases of ambiguity, the predicted protein sequence was analyzed using Interproscan (http://www.ebi.ac.uk/Tools/InterProScan/) for final domain prediction and validation of TF family.

Creation and Validation of the qRT-PCR Primer Set Library
To accurately quantify the expression levels of the identified putative regulatory genes, specific qRT-PCR primers were designed from each of the putative
regulatory gene sequences using PRIMEGENS (Xu et al., 2002). The specificity of each primer set was validated based on the number of amplicons produced after RT-PCR and qRT-PCR (i.e. only one amplicon was expected per RT/qRT-PCR at the average size of 100 bp; Supplemental Fig. S1, A and B). Moreover, 96 different amplicons were randomly selected for sequencing. This analysis returned 90 correct sequences out of the 96 analyzed (data not shown). Despite resequencing the last six amplicons, the sequences were of insufficient quality and, therefore, were discarded without a conclusion as to their identity. These data suggested that at least 94% of the primer sets are specific for the targeted putative regulatory genes. Since the 96 amplicons were randomly selected for sequencing, similar results should be expected for the entire primer library. Altogether, 1,280 primer sets were validated by these approaches.

The lack of specificity of the other primer sets is likely the result of the large number of soybean paralogs that arose from the whole genome duplications detected from analysis of the soybean genome (Schlueter et al., 2004, 2007). Although PRIMEGENS attempts to compensate for these effects, genome duplication is an additional difficulty in the design of specific primer sets against soybean transcripts. Based on the low divergence of homeologous genes, we assumed that amplicon size and dissociation curve profiles might not be sufficiently stringent to validate the primer set specificity. Accordingly, to confirm primer set specificity, we took advantage of the recent release of the soybean genome sequence (http://www.phytozome.net/soybean) to BLAST the 1,280 qRT-PCR primer sets. Such analysis also allowed us to attribute a soybean gene annotation number to each of the specific qRT-PCR primer sets. A total of 131 primer sets matched against two or three different soybean transcripts. Such primer sets were discarded, finally leading to the selection of 1,149 primer sets in the present soybean qRT-PCR library (Supplemental Table S2). These 1,149 primer sets allowed the quantification of 1,034 soybean genes (Fig. 1A).

In addition to their specificity, primer efficiency is another important characteristic to validate suitable qRT-PCR primer sets. Theoretically, under ideal conditions, the number of amplicons should increase by two after each PCR cycle. Based on an earlier study (Libault et al., 2007), only a few primer sets are highly efficient across a set of tested conditions. In this study, the efficiency of each putative soybean regulatory gene primer set was measured using the LinRegPCR software (Ramakers et al., 2003). The average primer set efficiency was calculated when gene expression was detected in at least 20 out of the 24 different cDNA samples tested. Out of the 1,149 primer sets used, 1,043 were retained and exhibited an average primer efficiency of 1.97 ± 0.05 (Supplemental Table S3). None of the primer sets displayed an efficiency of less than 60%, and 953 primer sets (91.4% of the library) had an efficiency of greater than 90% (Supplemental Fig. 1C).

These data provide confidence that the primer set library is of high specificity and can be used to accurately quantify putative regulatory gene mRNA expression levels.

Expression Pattern of Putative Soybean Regulatory Genes during Root and Nodule Development

The putative regulatory gene primer set library was first used to profile gene expression during nodule development. To identify regulated genes during nodulation, we compared gene expression in soybean roots inoculated with *Bradyrhizobium japonicum* and mock-inoculated roots at different stages of nodule development (4, 8, and 24 d after inoculation [DAI]). In our conditions, at 4 DAI, nodules were not visible but extensive root hair cell deformation could be seen, indicative of a strong plant response to *B. japonicum*. At 8 DAI, nodule primordia emerged, while nodules were fully developed at 24 DAI (i.e. leghemoglobin was present in the nodule [pink-red color], and nitrogen fixation occurred based on the absence of leaf chlorosis). Overall analysis of the expression pattern of soybean regulatory genes in mock-inoculated roots highlighted strong correlations between 4 and 8 DAI.
A lower correlation between 4- to 8-DAI and 24-DAI mock-inoculated roots suggested that gene expression was modified during root development. This analysis suggests that the regulatory gene expression profiles were well established and stably maintained during the first stages of root development.

During nodulation, the putative soybean regulatory gene expression profile was not affected during the first days following root inoculation by B. japonicum (i.e., the Euclidian distance between 4- and 8-DAI inoculated and mock-inoculated roots was low) but was strongly disrupted at 24 DAI (Fig. 1B). Such results suggested that a significant number of soybean regulatory genes were specifically expressed in mature nodules but not in developing nodules. On the other hand, based on our experience, mature nodules are plant organs extremely rich in mRNA. Consequently, comparisons between inoculated and mock-inoculated roots between 4 to 8 DAI and 24 DAI might also be explained by a strong dilution of the 4- to 8-DAI nodule mRNA by the overall root system mRNA, a dilution that was limited at 24 DAI when individual nodules can be harvested for analysis. These observations support the notion that a large number of putative regulatory genes were differentially expressed at 24 DAI.

We identified for each time point the differentially expressed putative soybean regulatory genes. For example, 126 soybean regulatory genes were differentially expressed during soybean nodule development (P < 0.05 and fold change < 0.5 or > 2; Supplemental Table S4). Among them, 31% (39 genes) were up-regulated, while 69% (87 genes) were down-regulated (Fig. 2). As expected based on the calculation of the correlation between the different conditions tested, most of the putative regulatory genes were differentially expressed at 24 DAI (five, 19, and 111 genes were regulated at 4, 8, and 24 DAI, respectively; Fig. 2). Among the 126 genes, only eight genes were found to be differentially regulated at multiple time points of the time course: three CCAAT-Box, one AP2/EREBP, one Myb, one C2H2 Zinc Finger domain, one bHLH, and one RNA-dependent RNA polymerase gene. All eight of these genes were exclusively up-regulated. Such genes might have an important role in multiple stages of nodule development (e.g., infection of plant cells by bacteroids or control of nodule development).

Characterization of the Roles of Putative Regulatory Genes in Nitrogen Assimilation/Metabolism and Nodule Development

A difficulty in profiling gene expression is choosing which tissues to compare. For example, during our analysis, it appears that 24-d mock-inoculated roots, in addition to lacking nodules, also exhibited symptoms of nitrogen starvation, such as chlorosis and leaf senescence. This stress condition might affect the expression pattern of the putative soybean regulatory genes, especially with respect to the comparisons between mock-inoculated and inoculated plants. Consequently, at 24 DAI, we made the assumption that some of the regulatory genes were regulated in response to nitrogen starvation in the mock-inoculated plants and did not have a specific role in nodule development. In order to obtain a clearer view of those genes that may be responding to the nitrogen provided by actively fixing bacteroids versus those that are strictly involved in the nodulation response, 24-DAI gene expression (inoculated versus mock inoculated) was compared in roots grown with and without nitrogen (KNO₃ versus KCl; Supplemental Table S4).

Of the 111 genes regulated in 24-DAI nodulated roots, only 12 were also responsive to root nitrogen status (1.2% of the 1,034 genes analyzed; Supplemental Table S4). Among these 12 genes, 10 were repressed during both nodulation and KNO₃ treatments. The remaining two genes were strongly up-regulated at 24 DAI but repressed in roots treated with KNO₃. The remaining 99 genes (9.6% Supplemental Table S4) were regulated specifically in nodulated roots, thereby supporting a role of these putative regulatory genes in the maintenance and activity of the nodule. The com-

**Figure 2.** Identification of putative soybean regulatory genes expressed differentially during nodulation. Venn diagrams showing the number of soybean genes significantly up-regulated (left) and down-regulated (right) during nodulation at 4 (yellow), 8 (red), and 24 (blue) d after B. japonicum inoculation. This analysis was performed on 1,043 different putative soybean regulatory genes. Genes regulated during nodulation were validated according to their fold change between inoculated (nodulated) and mock inoculated (nodule-free) roots (fold change > 2 or < 0.5). Student’s t test (P < 0.05) was performed between the three independent biological materials to ensure the significance of the differences.
comparison of KNO$_3$- and KCl-treated roots identified 38 genes responsive to nitrogen status but not involved in the nodulation process (3.7%; Supplemental Table S4). Recently, Ruffel et al. (2008) also studied the response of *M. truncatula* roots to different nitrogen treatments. In that study, plants were grown with or without 1 mM KNO$_3$ as well as under conditions where atmospheric nitrogen was present or absent in the environment of nodulated roots. Among the 752 TF genes analyzed, Ruffel et al. (2008) identified 101 TF genes regulated specifically in response to KNO$_3$ treatment (13.4%), 37 differentially expressed TF genes in response to the presence of atmospheric nitrogen (4.9%), and only 11 TF genes differentially regulated in response to both treatments (1.5%). This distribution was very different from our results with soybean. Such differences might be explained by the fact that in both studies the qRT-PCR primer libraries used only sampled a subset of the total TF genes that may be responding to nitrogen status. Of course, the data may also suggest fundamental differences by which these two plant species respond to nitrogen status. Finally, the experimental designs used by Ruffel et al. (2008) and in our experiments are also quite different, which could also explain the differences in the magnitude of the responses seen. While we compared *B. japonicum* inoculated with mock-inoculated roots, Ruffel et al. (2008) compared the expression patterns of TF genes on nodulated roots with or without atmospheric nitrogen.

**Putative Soybean Orthologs of *M. truncatula* and *L. japonicus* Nodulation-Related Genes Are Regulated during Soybean Nodulation**

We next compared the sequences of the 126 genes that we identified as differentially regulated in soybean nodule development with sequences of TFs from *L. japonicus* and *M. truncatula* with known roles in nodule development. Among them, Glyma09g10240 and Glyma10g10240 are homologous to *M. truncatula* HAP2.1 and its *L. japonicus* homologs CBF-A01 and CBF-A22 (Asamizu et al., 2008). To enlarge our investigation to other soybean genes sharing sequence similarity to *MtHAP2.1*, we BLAST searched the *MtHAP2.1* protein sequence against soybean genomic sequences. Based on our search criteria (e-value < $10^{-12}$ and score > 100), we identified four soybean homologous genes (Glyma09g36140, Glyma02g35190, Glyma19g38800, and Glyma10g10240). These same four genes were also identified by BLAST searching LjCBF-A01 and LjCBF-A22 protein sequences. To support the putative orthology between *MtHAP2.1* and these four soybean genes, we examined the syntenic relationship based on gene content, order, and orientation in the corresponding genomic regions, as previously performed by Zhang et al. (2007). Despite chromosome rearrangements (e.g., inversion of gene orientation), gene content was similar between the different chromosome regions analyzed, supporting that these genes are likely orthologous (Supplemental Fig. S2). While LjCBF-A22 was slightly induced several hours after bacterial inoculation, *MtHAP2.1* and LjCBF-A01 expression was strongly induced in nodules (Combier et al., 2006; Asamizu et al., 2008). Among the four soybean genes identified, Glyma10g10240 was strongly and consistently up-regulated during soybean nodulation, while Glyma02g35190 expression was significantly induced during the earlier steps of nodule development (Supplemental Table S4). Glyma03g36140 and Glyma19g38800 were not represented in the current qRT-PCR primer library. However, based on Illumina-Solexa sequencing of soybean cDNA, we confirmed that both genes were preferentially expressed in soybean root tissues including nodules (Supplemental Table S5).

In addition to the identification of the four putative soybean orthologs of *MtHAP2.1*, we also identified several soybean putative orthologs of *MtERN* (Glyma16g04410, Glyma16g27040, Glyma19g29000, Glyma02g08020, and Glyma08g12130). Based on this study, the expression of Glyma16g04410 was induced during early nodulation (Supplemental Table S4). An examination of the genomic regions of *MtERN* and Glyma16g04410 supports the notion that these genes are likely orthologous (Supplemental Fig. S3). A similar approach also identified Glyma19g29000 and Glyma08g12130 as putative *MtERN* orthologs. However, the genomic regions of Glyma16g27040 and Glyma02g08020 genes showed no microsynteny to the *MtERN* genome locus (Supplemental Fig. S3). Glyma16g27040, Glyma19g29000, Glyma02g08020, and Glyma08g12130 were not represented in the current qRT-PCR primer library. However, Glyma19g29000 and Glyma02g08020 were strongly expressed in soybean nodules, while Glyma16g27040 and Glyma08g12130 were not (Supplemental Table S5). Such observations reinforced the putative orthologous relationship between *MtERN*, Glyma16g04410, and Glyma19g29000. Note that even though the genome regions of Glyma08g12130 and *MtERN* showed significant microsynteny, Glyma08g12130 was not induced during nodulation. This result suggests that the regulation of Glyma08g12130 likely diverged from its progenitor following the duplication that gave rise to this paralog.

**The Expression of Soybean Homeologous Regulatory Genes during Nodulation**

The identification of several putative soybean orthologs of *MtHAP2.1* and *MtERN* is likely the result of the whole genome duplication events that occurred in soybean (Schlueter et al., 2004, 2007). The paralogs that arose from these duplications may show functional redundancy, which, among other issues, could present challenges to efforts to silence specific genes using RNA interference (RNAi) approaches in order to study nodulation phenotypes. A key question is whether these various paralogs show similar expression patterns suggestive of functional redundancy. To address
this in a more general way, we identified 46 pairs of soybean homeologous genes (92 genes) represented in the qRT-PCR primer library (M. Libault, A. Farmer, G.D. May, and G. Stacey, unpublished data). Among them, we identified four pairs with both putative regulatory genes regulated during nodulation, eight pairs with one of the two homeologs regulated during nodulation, and finally 34 pairs of putative regulatory genes without any of the two homeologous genes regulated during nodulation or in response to KNO3 treatment. Looking at their expression patterns, the four pairs responding to inoculation shared very similar expression patterns and also similar expression levels, with the exception of Glyma03g29190 and Glyma19g31940 (Fig. 3). With regard to the eight pairs where only one of the two genes was regulated during nodulation (Fig. 3), the data show a significant divergence in the expression profiles of the various paralogs (e.g. little or no expression of Glyma07g02630, Glyma18g04250, Glyma08g09970, and Glyma15g12930, while Glyma08g23380, Glyma11g34050, Glyma05g26990, and Glyma09g02030 were expressed at stronger levels; Fig. 3). In rare cases, the two homeologous genes shared very similar expression patterns and also similar expression levels, with the exception of Glyma03g29190 and Glyma19g31940 (Fig. 3). In a previous study, Koltai et al. investigated their expression patterns in a variety of soybean tissues. Interestingly, despite similar expression in response to inoculation. The identification of putative MtHAP2.1 and MtpERN soybean orthologous genes (Glyma02g35190, Glyma16g04410, and Glyma10g10240) induced during nodulation provides some validation to this approach. Among the different genes regulated, Glyma03g31980 and Glyma17g07330, two Myb TF genes, were found to be regulated at several time points of the time course (Fig. 4, F and K; Supplemental Table S6).

Based on further characterization (see below), Glyma03g31980 was named CND. The expression of CND and Glyma17g07330 during nodulation were similar. In a previous study, Koltai et al. (2001) reported the specific expression of the M. truncatula PHANTASTICA gene encoding a Myb TF in nodulated roots. Comparison of the M. truncatula PHANTASTICA protein sequence with that of the soybean CND and Glyma17g07330 MYB proteins using EMBOSs Pairwise Alignment Algorithms (http://www.ebi.ac.uk/Tools/emboss/align/) showed low degrees of identity and similarity (23.3% and 33.7% with CND and 20.3% and 34% with Glyma17g07330, respectively). In addition, no microsynteny exists between these three genes (data not shown). Therefore, to better characterize the role of these two TF genes, we investigated their expression patterns in a variety of soybean tissues. Interestingly, despite similar expression during nodule development, CND and Glyma17g07330 expression showed tissue-specific differences. Whereas CND was mainly expressed in root hair cells and the shoot apical meristem, the expression of Glyma17g07330 was very low in most of the tissues tested, with the exception of striped root (i.e. root devoid of root hair cells) and root tissues (Supplemental Fig. S5).

To further examine the expression of CND during nodule development, promoter-GUS and -GFP fusions were constructed and introduced into soybean roots
by *Agrobacterium rhizogenes* hairy root transformation. A 2-kb region 5' of the CND gene was identified from the Phytozome Web site and cloned upstream of the GFP and GUS reporter genes to construct transcriptional fusions. Early during nodule development (13 DAI), transformed roots showed CND promoter-GFP

**Figure 3.** Expression of soybean homeologous genes during nodulation and in response to KNO₃ and KCl treatments. The expression of each soybean gene (black bars) and its respective homeolog (gray bars) was quantified by qRT-PCR after normalization with the soybean reference gene *Cons6* (y axis). Eight different conditions were used to quantify expression (4, 8, and 24 DAI inoculated [IN] and mock inoculated [UN] and in response to KNO₃ and KCl treatments; x axis). These 12 pairs of homeologous genes were divided in two subgroups (A–D, both homeologs were significantly regulated during nodulation; E–L, only one of the two homeologs was significantly regulated during nodulation; for details, see Supplemental Table S2).
expression in emerging nodules and, at a lower level, in root cells (Fig. 5, A–D). Although CND expression was apparent throughout the root, GFP signal was noticeably absent in emergent secondary root tips (Fig. 5, A and B). This result was also supported by the analysis of roots expressing the CND promoter-GUS construct, although in rare occasions secondary root tips did show some GUS staining (Fig. 5C, white arrow). CND expression was induced very early during nodule primordium formation, at the location of the infected zone (Fig. 5, C and H, black arrows). Later during nodule development (28 DAI and later), CND expression decreased in the infection zone of the nodule (Fig. 5, E, F, and I). Looking at transverse sections of GUS-stained transgenic roots and developing and mature nodules, CND was strongly expressed in the vascular tissues of all organs (Fig. 5, G–I). At a finer level, GUS staining was also observed in epidermal, endodermal, and cortical cells. These observations support a change in the expression pattern of CND during nodule development, especially in the infected zone of the nodule.

Characterization of the Role of CND during Nodule Development

RNAi silencing is a common strategy to characterize gene function when mutants are not available. Several studies focusing on legume nodulation have successfully used this strategy to investigate gene function. For example, silencing of \( \text{MtDMI2} \) led to 75% of roots with no nodules compared with 0% to 20% of control roots with no nodulation (Limpens et al., 2003). Using a similar RNAi strategy, Vernié et al. (2008) showed an increase in nodule formation and infection threads after rhizobial inoculation coincident with the silencing of \( \text{MtEFD} \), a \( \text{MtERF} \) TF gene regulated during \( M. \ truncatula \) nodulation (El Yahyaoui et al., 2004). However, the authors concluded that silencing of \( \text{MtEFD} \) created a weak allele relative to the \( \text{efd} \) null mutant. In
addition, the silencing of MtCDPK1 led to smaller roots and decreases in nodule formation and mycorrhizal infection (Ivashuta et al., 2005).

Because the transcriptional response of a gene to a specific treatment does not necessarily imply an essential role in this process, we further investigated the role of CND in nodule formation. To specifically silence CND, a DNA fragment corresponding to the predicted 3’ untranslated region (UTR) was cloned to create a hairpin structure (see “Materials and Methods”). As a control, pCGT5200 carrying an RNAi construct specific to the GUS gene was used. Three independent experiments were performed allowing the counting of nodule numbers on 44 CND-RNAi and 36 control RNAi-GUS roots. The silencing of CND consistently affected nodule development by significantly reducing nodule numbers by approximately 40% compared with the RNAi-GUS control roots (Student’s t test, P = 0.03; Fig. 6A). There was also a slight increase in nodule size that coincided with the drop in nodule numbers in the CND-silenced roots (i.e. nodule size increased by around 20% in CND-RNAi roots when compared with the control (Student’s t test, P < 0.001; Fig. 6B). A relationship between nodule size and nodule number was previously investigated in the supernodulating soybean mutant nts382 and its wild-type parent (soybean cv Bragg; Day et al., 1987). In this case, more nodules in the mutant roots resulted in a smaller average nodule size. Similarly, the hypernodulation mutant Ljtml also exhibited smaller nodules (Magori et al., 2009). In keeping with this relationship, the increase in nodule size in the CND-RNAi-silenced roots is likely an indirect effect of the total decrease in nodule numbers.

Based on the expression of CND in the infected zone of young nodules, we investigated more closely the infection of CND-RNAi nodules by B. japonicum. CND-RNAi and RNAi-GUS nodules were stained with SYTO-13, a nucleic acid-binding dye (Veereshlingam et al., 2004). By laser-scanning confocal microscopy, we did not observe a significant modification in the pattern of infected cells coincident to the silencing of the CND gene (Supplemental Fig. S6). Both the CND-RNAi nodules and the control nodules exhibited a clearly distinguishable infected zone.

To confirm the silencing of CND, we quantified gene expression in RNAi-GUS control and CND-RNAi isolated nodules using qRT-PCR (i.e. for each of the three
biological replicates of CND-RNAi and RNAi-GUS roots, small, medium, and large nodules were pooled and RNA was extracted). These experiments confirmed the silencing of CND at all stages of nodule development (Fig. 6C). We assume that the fact that nodule numbers were only reduced by 40% is likely due to the fact that 100% silencing of CND gene expression was not achieved. We also cannot rule out the possibility of functionally redundant TF genes that were not silenced by the CND-RNAi construct.

Although primers were designed to specifically silence CND expression, the similarity of sequences between the members of the Myb TF family may have resulted in silencing of more than one gene. To test this hypothesis, the CND nucleotide sequence was BLAST searched against the available soybean genomic sequence (http://www.phytozome.net/soybean.php). The closest homolog was Glyma19g34740 (score = 379, e-value = 6.2e⁻¹⁰³). Based on the strong identity between CND and Glyma19g34740 (83% and 87% identity of their nucleotide and amino acid sequences, respectively), we hypothesized that Glyma19g34740 might be homeologous to CND and that the function of these two Myb genes might be redundant. To better establish the relationship between these two genes, we mined soybean genome sequences to identify genes surrounding both Myb genes. Not surprisingly, the environment of these two genes in terms of gene content, order, and orientation was very similar, supporting their homeology (Supplemental Fig. S7). The strong nucleotide sequence identity between CND and Glyma19g34740 suggested that both genes might be silenced during our experiments. Accordingly, we also quantified the expression levels of Glyma19g34740 in the CND-RNAi and RNAi-GUS control nodules. The data showed that our construct specifically silenced the expression of the CND Myb gene (Fig. 6D). In addition, these analyses also demonstrated the strong expression of Glyma19g34740 in nodules, supporting a possible partial redundancy in the function of these two Myb genes. This result might be a second factor explaining the mild phenotype of the CND-RNAi plants.

CONCLUSION

The development and use of our soybean qRT-PCR library allowed quantification of the expression of 1,034 soybean putative regulatory genes. By focusing on nodule development, we validated the use of this approach by identifying putative soybean orthologs of MtHAP2.1 and MtERN as regulated during nodulation. In addition, using RNAi silencing, we identified a Myb TF gene, CND, involved in nodule formation. In
addition to PHANTASTICA, our data suggest that probably several Myb genes including CND might be involved in nodule development.

MATERIALS AND METHODS

Bacterial Culture

DHsEs Escherichia coli and K999 Agrobacterium rhizogenes were grown in Luria-Bertani medium supplemented with appropriate antibiotics at 37°C and 30°C, respectively. Bradyrhizobium japonicum USDA110 was grown at 30°C for 3 d in HM medium (Cole and Elkan, 1973) supplemented with yeast extract (0.025%), d-Ara (0.1%), and chloramphenicol (0.004%). Before plant inoculation, B. japonicum cells were pelleted (4,000 rpm for 10 min), washed, and dialyzed against the Phytorector buffer appropriate for concentration according to the experiments (i.e. optical density at 600 nm [OD600] = 0.1 or 0.3 when inoculated with wild-type soybean [Glycine max] seedlings and hairy roots, respectively).

Cloning

To silence the CND Myb gene, a 120-bp fragment specific to the 3’ UTR of GmMyb was amplified from soybean cDNA using the following primers: GmMyb-RNAdifor (5’-TTGAGTTACACGCTTAAATTGACA-3’) and GmMyb-RNAdifors (5’-GACCAAGCTTCTATTCAAGCA-G’). The antisense fragment was cloned into the entry vector CTG11050 by TA cloning using two AphII sites (engineered to produce 3’ T overhangs). The RNAi entry vector was recombined with the destination vector CTG11017A (Supplemental Fig. S8) using the LR Clonase reaction (Gateway LR Clonase II enzyme mix; Invitrogen). The LR Clonase reaction combining the gene fragment in the entry vector and destination vector CTG11017A was used to produce GmCND-RNAi vector. The RNAi control vector used in these studies (CGT3200) was previously described by Govindarajulu et al. (2009). The fidelity of the clones was verified by sequencing, and the clones were electroporated into A. rhizogenes strain K999 and used for composite plant production (see below).

The Gateway system (Invitrogen) was used to clone GmMyb promoter upstream to the GFP and GUS cDNAs. By mining genomic sequences available on the Phytorector website (http://www.phytorector.net/soybean.php), we identified a 1,967-bp DNA fragment 5’ to the first codon of the CND gene. By two independent PCR, we created the AttB sites at the extremities of the promoter sequences. We used soybean cv Williams 82 genomic DNA as template and the following primers: GmMybpromotorAttFor (5’-AAAAAG-CAGGCTCCTACATATGGTGATTTCGTTAATATTAAATAA-3’) and GmMybpromotorAttRev (5’-AAGAAGATTGTTTCTGATACTAGTACCTCCCTC) first and the universal AttB primers attB1adaptor (5’-CAGGCAACATTCTGACAA-3’) and attB2adaptor (5’-GGGGACCATTTGTACAGAAAAGTCTCGTG-3’) second.

Using the Gateway BP Clonase II enzyme mix, the GmMyb promoter fragment was introduced first into the pDONR-Zeo vector (Invitrogen) and then into pX1T or pX2T destination vector using the Gateway LR Clonase II enzyme mix (Invitrogen). The pX1T and pX2T destination vectors carry the GUS and GFP reporter genes, respectively (Xiao et al., 2005).

A. rhizogenes (strain K999) was transformed by electroporation with pCGT11017-3’UTR-GmMyb, GM Mybpromotor-pX1T, and GM Mybpromoter-pX2T vectors.

Plant Culture

Soybean seeds were surface sterilized according to Wan et al. (2005) and then placed on moist filter paper for germination in dark conditions, 80% humidity, and 27°C for 3 days. The three-day-old seedlings were grown on moist filter paper and transferred into vermiculite-perlite (3:1) supplied with B&D medium (Broughton and Dilworth, 1971), while nodulated or nodule-free roots produced to quantify gene expression during nodule development were watered with B&D medium not supplemented. Seedlings were inoculated with 1 mL of B. japonicum suspension in water (OD600 = 0.1) or only water (mock). B&D medium was supplemented with KNO3 or KCl (10 mM final concentration) when analyzing putative regulatory gene expression in response to nitrogen starvation. Nitrogen (KNO3) and nitrogen-free (KCl) roots, nodulated or nodule free, were harvested 24 d after treatment and 0, 4, 8, 16, 24, and 32 DAI. In all cases, root apices (around 1 cm) were removed prior to freezing in liquid nitrogen. Two to six plants were harvested by biological replicates. These experiments were repeated at least three times on different set of plants to ensure reproducibility in the plant tissues analyzed.

Soybean Regulatory Gene Expression Profile during Nodulation

Soybean hairy root transformation was done essentially as described by Taylor et al. (2006). Two-week-old soybean shoots were cut between the first true leaves and the first trifoliate leaf and placed into rock-wall cubes (Fibrgro). Each shoot was inoculated with 4 mL of A. rhizogenes (OD600 = 0.3) and then allowed to dry for approximately 3 d (23°C, 80% humidity, long-day conditions) before watering with deionized water. After 1 week, the plants were then transferred to pots with vermiculite/perlite mix (3:1) wetted with nitrogen-free plant nutrient solution (Lullien et al., 1987). One week later, the shoots were transferred to the greenhouse (27°C, 20% humidity, long-day conditions). Two weeks after vermiculite-perlite transfer, the shoots were inoculated with B. japonicum (10 mL, OD600 = 0.08).

CND-RNAi and control nodulated transgenic roots were isolated (these roots expressed the GFP reporter gene), and the nodules were isolated and then categorized based on the presence of leghemoglobin (a pink-red color in the infected zone of the nodule was used as a marker of nodule maturity; leghemoglobin gene expression increases proportionally with the age of the nodule [Marcker et al., 1984], and its induction occurs before the detection of nitrogenase activity [Verma et al., 1979]) and nodule size (i.e. leghemoglobin-free nodules and small [<3.5 mm2], medium [3.5 mm2 and <4.5 mm2], and large [>4.5 mm2] mature nodules with leghemoglobin).

For each category, nodule numbers were counted with a Leica MZFLIII stereomicroscope equipped with epifluorescence excitation and a GFP long-pass filter (500 nm long pass; no. 41018; Chroma Technology).

Molecular Devices MetaMorph 7.5 was used to measure the area of each nodule (the areas of 73 large, 182 medium, 109 small mature, and 33 immature nodules from RNAi-GUS and 54 large, 178 small, 88 small mature and 32 immature nodules from CND-RNAi were measured).

To observe B. japonicum infection in transgenic nodules, CND-RNAi and RNAi-GUS nodules were embedded in paraffin before sectioning. Five-micrometer sections were stained during 20 min with 10 μM SYTO13 in 1 M sodium phosphate buffer (pH 7.0). Stained sections were observed with a Zeiss LSM 510 META confocal microscope (488 nm excitation wavelength and 500- to 550-nm emission filter).

GmMyb promoter-GUS and -GFP transgenic roots were observed 13, 18, 29, and 40 DAI. GmMyb promoter-GUS roots were fixed and stained as described by Govindarajulu et al. (2008). To section roots and nodules stained with GUS, 80-μm-thick sections were obtained using a fully automated Vibratome 3000 Plus after embedding the tissues in 6% low-melting agarose. Roots with GUS-stained nodules were observed using a SMZ1500 Nikon microscope equipped with a DXM1200 Nikon digital camera, while GmFW115 promoter-GFP roots were observed using a Leica MZFLIII stereomicroscope equipped with epifluorescence excitation and a GFP long-pass filter (500 nm long pass; no. 41018; Chroma Technology).

Illumina-Solexa Sample Preparation, Read Alignment, and Statistical Analysis

mRNAs isolated from different soybean tissues were reverse transcribed using standard protocols. After first- and second-strand cDNA synthesis, the cDNAs were end repaired prior to ligation of Soxela adaptors. The products were sequenced on an Illumina-Solexa platform.

Illumina Genome Analyzer II image data were base called and quality filtered using the default filtering parameters of the Illumina GA Pipeline GERALD stage. Alignments of passing 36-mer reads to all contigs of the Glyma1 8% Soybean Genome assembly (Soysbean Genome Project, Department of Energy Joint Genome Institute) were performed using GSNAp, an alignment program derived from GMAP (Wu and Watanabe, 2005) with optimizations for aligning short transcript reads from next-generation sequencers to genomic reference sequences. Alignments were processed using the Illumina pipeline (Bol et al., 2008), keeping only alignments that had at least 34 out of 36 identities and had no more than five equivalent best hits. Read counts used in expression analyses were based on the subset of uniquely aligned reads that also overlapped the genomic spans of the Glyma1 gene predictions.
qRT-PCR Primer Design

To ensure maximum specificity of the designed primers, we used PRIME-GENS (http://digbio.missouri.edu/primersgen/; Xu et al., 2002). This software compares the sequence of each primer designed against a database of the 2,258 putative soybean regulatory genes identified in this study as well as the G. max Unigene database to ensure that the designed primers are unique to the sequence and are likely to give specific amplification. The following criteria were used to design the primers: melting temperature of 60°C, PCR amplicon lengths of 80 to 120 bp, and yielding primer sequences with lengths of 19 to 23 nucleotides with an optimum at 21 nucleotides and GC contents of 40% to 60%. The list of the validated primers described in this study is available in Supplemental Table S2.

To quantitatively express the expression of the Glyma19g34740 gene, Glyma19g34740-forward (5'-CAAAGACCAAAACTAGAACA-3') and Glyma19g34740-rev (5'-TGGTAGACAACAAAGGAGGA-3') primers were designed. Primer specificity was confirmed by dissociation curve profile after qRT-PCR amplification.

RNA Extraction, DNase Treatments, and RT

Total RNA was isolated using Trizol Reagent (Invitrogen) according to the manufacturer’s instructions followed by chloroform extraction to improve their purity. Purified RNA was treated with TURBO DNase (Ambion) to remove any contaminating genomic DNA according to the manufacturer’s instructions. In the experiments, 100 μg of DNA-free total RNA was used for first-strand cDNA synthesis using Moloney murine leukemia virus (Promega) according to the manufacturer’s instructions. The lack of genomic DNA contamination was verified by qRT-PCR using primers designed against soybean genomic DNA. Moreover, cDNA synthesis efficiency was analyzed by amplifying by qRT-PCR Const6 cDNA with two different primer sets separated by 1 kb (Const6-forward, 5'-AGATGGAAATCTCGAGGT-3', and reverse, 5'-CTAATGGCAATTGCAGCTCTC-3'; Const5-forward, 5'-AAAGTGAAATTCGCTTTCC-3', and reverse, 5'-CCGAAATCTGCGCAATACTGA-3'). For each condition and replicate tested, less than one PCR cycle difference was measured between both primer sets, supporting the high quality of the cDNA synthesis.

qRT-PCR Conditions and Data Analysis

qRT-PCR was performed as described by Libault et al. (2008). Primer set specificity was confirmed by analyzing the dissociation curve profile of each qRT-PCR amplicon, and the efficiency of primers (Peff) was quantified using LinRegPCR (Ramakers et al., 2003).

Validation of the specificity and efficiency of the qRT-PCR amplicon, and the efficiency of primers (Peff) was quantified using LinRegPCR (Ramakers et al., 2003). The cycle threshold (Ct) value of the reference gene was separated by 1 kb (Const6-3 and Const5-3). For each condition and replicate tested, less than one PCR cycle difference was measured between both primer sets, supporting the high quality of the cDNA synthesis.

Supplemental Data

The following materials are available in the online version of this article.

Supplemental Figure S1. Validation of the specificity and efficiency of the qRT-PCR primers designed to quantitate putative soybean regulatory genes.

Supplemental Figure S2. Syntenic relationship between putative soybean orthologous genes to MtHAP2-1 and MtHAP2-1.

Supplemental Figure S3. Syntenic relationship between putative soybean orthologous genes to MiERN and MiERN.

Supplemental Figure S4. Expression of soybean homeologous genes during nodulation and in response to KNO3 and KCl treatments.

Supplemental Figure S5. Expression patterns of Glyma03g31980 (CND) and Glyma17g07330 in 10 soybean tissues.

Supplemental Figure S6. Confocal light micrographs of CND-RNAi nodule (left) and RNAi-GUS (right) sections.

Supplemental Figure S7. Syntenic relationship between Glyma03g31980 and Glyma19g34740.

Supplemental Figure S8. Construction of the RNAi vector CGTI1017A.

Supplemental Table S1. Identification of putative soybean regulatory genes from public cDNA databases.

Supplemental Table S2. Gene expression analysis for the 1,149 putative regulatory genes during nodulation and in response to KNO3.

Supplemental Table S3. Quantification of primer set efficiency in a large number of independent qRT-PCRs.

Supplemental Table S4. Expression of 162 soybean putative regulatory genes during nodulation and in response to KNO3, and KCl treatments.

Supplemental Table S5. Expression of putative soybean orthologs of MhHAP2.1 and MiERN genes in different soybean tissues.

Supplemental Table S6. Time course of expression of the putative soybean regulatory genes during nodulation (i.e. 0–32 d after B. japonicum inoculation) or after the KNO3 treatment.

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