A rice calcium-dependent protein kinase is expressed in cortical root cells during the presymbiotic phase of the arbuscular mycorrhizal symbiosis

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Abstract

Background: The arbuscular mycorrhizal (AM) symbiosis consists of a mutualistic relationship between soil fungi and roots of most plant species. This association provides the arbuscular mycorrhizal fungus with sugars while the fungus improves the uptake of water and mineral nutrients in the host plant. Then, the establishment of the arbuscular mycorrhizal (AM) symbiosis requires the fine tuning of host gene expression for recognition and accommodation of the fungal symbiont. In plants, calcium plays a key role as second messenger during developmental processes and responses to environmental stimuli. Even though calcium transients are known to occur in host cells during the AM symbiosis, the decoding of the calcium signal and the molecular events downstream are only poorly understood.

Results: The expression of seventeen Calcium-dependent Protein Kinase (CPK) genes representative of the four distinct phylogenetic groups of rice CPKs was monitored during the presymbiotic phase of the AM symbiosis. Among them, OsCPK18 and OsCPK4, were found to be transcriptionally activated in response to inoculation with the AM fungus Glomus intraradices. OsCPK18 and OsCPK4 gene expression was also up-regulated by fungal-produced diffusible molecules. Laser microdissection revealed expression of OsCPK18 in cortical cells, and not in epidermal cells of G. intraradices-inoculated rice roots, suggesting a preferential role of this gene in the root cortex. Moreover, a plasma membrane localization of OsCPK18 was observed by transient expression assays of green fluorescent protein-tagged OsCPK18 in onion epidermal cells. We also show that the myristoylation site of the OsCPK18 N-terminus is required for plasma membrane targeting.

Conclusion: The rapid activation of OsCPK18 expression in response to AM inoculation, its expression being also induced by fungal-secreted signals, together with the observed plasma membrane localization of OsCPK18, points to a role for OsCPK18 in perception of the AM fungus. The OsCPK18 gene might be considered as a marker for the presymbiotic phase of the symbiotic process. These findings provide a better understanding of the signaling mechanisms operating during the AM symbiosis and will greatly facilitate their molecular dissection.
Ca2+ spiking responses are elicited in though both cytoplasmic [10] and nuclear [9] pre-infection transcriptional reprogramming of the root [8-12]. Even cellular calcium, alterations in the cellular architecture and host cells triggers a rapid and transient elevation of intra-diverse plant species [7]. Perception of Myc factors by the charides which stimulate formation of AM symbiosis in reported that the AM fungus secretes lipochitooligosaccharides which are analogous to the rhizobial Nod factors). Very recently, it was turn, produce diffusible molecules, the "Myc factors" (analogous to the rhizobial Nod factors). Very recently, it was reported that the AM fungus secretes lipochitooligosaccharides which stimulate formation of AM symbiosis in diverse plant species [7]. Perception of Myc factors by the host cells triggers a rapid and transient elevation of intracellular calcium, alterations in the cellular architecture and transcriptional reprogramming of the root [8-12]. Even though both cytoplasmic [10] and nuclear [9] pre-infection Ca2+ spiking responses are elicited in M. truncatula roots in response to AM fungi, the mechanisms by which Ca2+ alterations are sensed and transduced into early AM-induced signaling remain unknown.

Once contact between the symbionts is established, the fungus enters into the root through the epidermal cells, and penetrates into the cortex where it forms highly branched structures, called arbuscules, in the cortical cells of the root. The arbuscules are the site of the major nutrient exchange between the two symbionts [2,13,14]. It is also known that the plant response to Myc factors is mediated by a partially characterized signaling pathway which is required for the establishment of both rhizobial and AM symbioses, the so called common symbiosis (SYM) pathway [2,13-15]. Forward genetic analysis in the model legumes Medicago truncatula and Lotus japonicus has led to the identification of components of the SYM signaling pathway. They are: a leucine-rich-repeat receptor-like kinase, the SYMRK protein in L. japonicus (known as DMI2 for "Does Not Make Infections 2" in M. truncatula), two nucleoporins (NUP85 and NUP133), two cation channel proteins (the L. japonicus CASTOR and POLLUX proteins; DMI1a and DMI1b in M. truncatula), a calcium and calmodulin-dependent protein kinase (CCaMK in L. japonicus; DMI3 in M. truncatula) and CYCLOPS (LjCYCLOPS; DIM3-interacting protein in M. truncatula) [16-21]. CCaMK interacts with, and phosphorylates, CYCLOPS in the nucleus [21,22]. In rice, the function of several SYM genes appears to be conserved, including CASTOR and POLLUX (acting upstream of the calcium-spiking signal) and CCaMK and CYCLOPS (acting downstream of the calcium-spiking signal) [23-25]. Evidence also support the existence of alternative, SYM-independent signaling pathways controlling the early responses to AM fungi in both rice and M. truncatula [25,26].

Transcript profiling of mycorrhizal roots allowed the identification of AM-regulated genes in several plant species, including rice [3,27-30]. However, the majority of these studies focused on the mature phase of the symbiotic process, a period in which the host root is already colonized and arbuscules are developed in the root cortical cells. Along with this, alterations in the expression of genes connected to nutrient acquisition processes, such as phosphate transporter genes, are well documented in different AM associations [31,32]. Genes involved in cellular modifications, transcriptional control and defense-related responses are also known to be regulated during the AM symbiosis [4,31].

Even though alterations in Ca2+ levels are known to occur in host cells during the presymbiotic phase, the decoding of the calcium signal is only poorly understood. On the other hand, it is well established that Calcium-dependent protein kinases (CPKs or CDPKs) are important Ca2+ sensors in signaling processes during growth, development and stress responses in plants [33,34]. CPKs belong to the CDPK/SnRK superfamily of protein kinases and represent a differentiated group of protein kinases found in plants, algae and protists [34-36]. They possess a characteristic structure consisting of four domains: an amino terminal variable domain, a serine/threonine kinase domain, a junction autoinhibitory domain, and a C-terminal calmodulin domain. These features make CPKs ideally structured to rapidly perceive alterations in intracellular calcium concentration and translating them into protein phosphorylation
cascade. CPK functioning is, however, different from that of CcAMK functioning, since CPKs do not require calmodulin for their activation. Ca<sup>2+</sup> binds directly to the calmodulin domain of CPKs and induces a conformational change resulting in kinase activation [34]. The available information on plant CPKs from various plant species indicates that they are encoded by multigene families and that whereas some of the genes are ubiquitously expressed, others show a tissue-specific pattern of expression or are regulated by stress (wounding, salinity, cold, drought, pathogen infection) [33,37,38].

Knowing that Ca<sup>2+</sup> plays a central role in the AM-induced signaling pathway, it was of interest to investigate to what extent CPKs are involved in the AM-induced signaling pathway. Towards this goal, the expression pattern of seventeen cpk genes was monitored in rice plants that have been inoculated with the AM fungus Glomus intraradices. We provide evidence that the expression of two distinct cpk genes, the OsCPK18 and OsCPK4 genes, is rapidly induced during the presymbiotic phase of the rice/G. intraradices interaction. OsCPK18 and OsCPK4 gene expression is also activated by fungal-produced diffusible fungal signal(s). By using the laser microdissection (LMD) technology, OsCPK18 expression was detected in cortical cells, but not epidermal cells, of the G. intraradices-inoculated rice roots. Moreover, a plasma membrane localization of OsCPK18 is here reported, the myristoylation site of OsCPK18 being required for its plasma membrane localization. Together, these findings suggest that OsCPK18 might play a role during recognition of the AM fungus by the host cells.

**Results**

**Expression of CPK genes in AM-inoculated rice roots**

A genome-wide analysis of rice CPK genes identified 31 genes which are distributed into four phylogenetic groups (I-IV) [39,40]. Moreover, a comparison of the rice CPK genes distinguished 11 closely related pairs which, most probably, have arisen via sequential duplication events, the OsCPK1/15, OsCPK2/14, OsCPK3/16, OsCPK4/18, OsCPK5/13, OsCPK7/23, OsCPK8/20, OsCPK11/17, OsCPK21/22, OsCPK24/28 and OsCPK25/26 pairs [40]. Based of the homology and phylogenetic relatedness among the rice CPK genes, we selected a subset of seventeen CPK genes representative of the four distinct phylogenetic groups of rice CPKs in which at least one representative member for each pair of closely related CPK genes was present. The subset of genes assayed in this work included OsCPK7, OsCPK10, OsCPK13, OsCPK17 and OsCPK24 from Group I; OsCPK2, OsCPK15, OsCPK19 and OsCPK25 from Group II; OsCPK8, OsCPK9, OsCPK16 and OsCPK22 from Group III; and OsCPK4, OsCPK18, OsCPK30 and OsCPK31 from Group IV.

The expression pattern of selected rice CPK genes and their transcriptional response to inoculation with the AM fungus G. intraradices, were examined during the presymbiotic phase of the symbiotic process. A preliminary screening was carried out by semiquantitative RT-PCR experiments with RNA samples obtained from whole rice roots that had been inoculated with fungal spores using the single sandwich method. Total RNA was isolated at 24, 48, 72 and 96 hours after inoculation of the rice roots with G. intraradices, as well as from mock-inoculated rice roots. Many CPK genes were found to be expressed in rice roots and at different levels (Additional file 1: Figure S1). Among them, the OsCPK4 and OsCPK18 genes were expressed at the highest levels. Moderate to low levels of expression were observed for OsCPK10, OsCPK13, OsCPK17, OsCPK24, OsCPK15, OsCPK19, OsCPK8 and OsCPK9, whereas OsCPK30 transcripts were barely detected (Additional file 1: Figure S1). The OsCPK7 and OsCPK16 genes showed expression profiles similar to those shown for OsCPK8 and OsCPK30, respectively (results not shown). Taken in the whole, the expression level of the various CPK genes here investigated appears not to be dramatically affected upon inoculation with G. intraradices, with the exception of OsCPK4 and OsCPK18 expression (Additional file 1: Figure S1).

For a comparison, the expression of the known SYM genes from rice, namely the OsSYM1, OsPOLLUX, OsCASTOR and OsCCaMK genes, was also examined. This analysis revealed up-regulation of OsSYM1, OsPOLLUX and OsCCaMK in response to G. intraradices at 72 and 96 hours post-inoculation (Additional file 1: Figure S1). The observed induction of these genes indicates that the host plant cells perceive and respond to the AM fungus through the activation of the AM-specific SYM signaling pathway.

Since RT-PCR analyses do not provide reliable quantitative data of gene expression, quantitative reverse transcription-PCR (RT-qPCR) was used to further characterize the effect of G. intraradices inoculation on OsCPK4 and OsCPK18 gene expression. By using the single sandwich system for fungal inoculation, up-regulation of OsCPK18 gene expression occurred as early as 24 h post-inoculation with G. intraradices (Figure 1A, upper panel). The level of OsCPK18 transcripts remained higher at the subsequent time points in the G. intraradices-inoculated roots compared to mock-inoculated roots. Concerning OsCPK4, its expression was also found to be up-regulated in response to G. intraradices inoculation during the time period of 24-72 hours. OsCPK4 expression returned to a level similar to that of non-inoculated roots by 96 h post-inoculation (Figure 1A, middle panel). Finally, OsCCaMK expression increased in G. intraradices-infected roots relative to mock-inoculated roots by 72-96 hours post-inoculation (Figure 1A, lower panel).
Overall, gene expression studies revealed up-regulation of the rice OsCPK4 and OsCPK18 genes during the presymbiotic phase of the AM symbiosis. These results were consistently observed in all three independent experiments. Although the expression of the OsCPK18, OsCPK4 and OsCCaMK genes was up-regulated in AM-inoculated roots compared to non-inoculated roots, it is also true that the amplitude of the differential expression for these genes was not very high during the time period here analyzed. Concerning OsCCaMK for which a role during AM symbiosis has been demonstrated in rice [23], its variation in the expression level in response to AM inoculation is also low and appears to occur at a later time point compared to the observed activation of OsCPK18 and OsCPK4 gene expression.

**Figure 1** Gene expression analysis by real-time qPCR for the OsCPK18, OsCPK4 and OsCCaMK genes. The single-sandwich (A) or the double-sandwich (B) system was used for inoculation. Roots inoculated with G. intraradices and mock-inoculated roots were harvested at the indicated times. Each sample consisted on a pool of at least 12 individual plants. Expression levels are shown relative to the housekeeping OsAct1 gene. Data shown represents the means ± error. Three independent experiments were carried out with similar results.
Diffusible factors released by *G. intraradices* induce *OsCPK18* and *OsCPK4* expression in rice roots

It is generally assumed that plants perceive AM fungi even before physical contact between the two symbionts, and that recognition of Myc factors triggers alterations in Ca$^{2+}$ levels and transcriptional responses in host roots [7,9,11,41]. In this work, the double sandwich method was used to investigate whether the observed induction of *OsCPK18* and *OsCPK4* expression is attributable to diffusible factors released by the fungus. This system prevents contact between the two symbionts while allowing the exchange of signal molecules [42]. *OsCPK18* and *OsCPK4* expression was analyzed by RT-qPCR (Figure 1B). When using the double sandwich system for inoculation of rice roots, *OsCPK18* and *OsCPK4* expression was found to be rapidly activated in response to *G. intraradices* inoculation (Figure 1B, upper and medium panel). However, induction of *OsCPK4* and *OsCPK18* expression was not maintained with time (the maximum induction occurred at 24 h post-inoculation for the two genes). Similar levels of transcript accumulation were observed in *G. intraradices*- and mock-inoculated roots at the latest time point here analyzed (96 hours post-inoculation). From these results it can be concluded that a diffusible fungal factor elicits expression of the rice *OsCPK18* and *OsCPK4* genes, and that this activation is transient. Most probably, contact between the two partners is needed to maintain the expression of these genes in an activated manner at the subsequent stages of the infection process. Under the same experimental conditions, an activation of *OsCCaMK* gene expression also occurred at 24 h post-inoculation although differences in *OsCCaMK* gene expression between AM-inoculated and mock inoculated roots were lower than those observed for the *CPK* genes (Figure 1B, lower panel).

**OsCPK18** expression in microdissected root cells

The laser-microdissection (LMD) technology has been successfully used for gene expression analysis in arbuscule-containing cells in different plant species such as *Medicago*, *Lotus* or tomato [27-29,43,44]. A variety of protocols have been developed for LMD of root tissues in order to identify the most appropriate fixation and embedding conditions that preserve cellular morphology, while still enabling extraction of high quality RNA for PCR amplification. In this way, laser microdissected cells can be used for RNA extraction and expression studies, thus avoiding the dilution effect of RNA samples extracted from whole roots. In this work, the protocol previously developed [43] for the isolation of cells from tomato roots was applied for the acquisition of rice root cells. The use of paraffin tissue preparations coupled to Methacarn fixation provided rice root tissues that satisfactorily retain the cellular morphology. Next, RNA samples of high quality were obtained from laser microdissected root cells.

Sections of the epidermis and the cortex were prepared from *G. intraradices*- and mock-inoculated rice roots at four days after inoculation (Figure 2). Cells, either epidermal or cortical, were collected pooled and used for RNA extraction. The cell type-specific pattern of expression of the *OsCPK18* gene was examined in laser microdissected cells. As it is shown in Figure 2F, *OsCPK18* transcripts were exclusively detected in cortical cells of *G. intraradices*-inoculated rice roots. *OsCPK18* transcripts were not detectable in epidermal cells of the fungal-inoculated roots. The absence of PCR amplification products in epidermal cells of the fungal-inoculated roots was confirmed by nested PCR (results not shown). Transcripts for the *ubiquitin1* gene (Figure 2F) or the *cyclophilin* gene (results not shown) were also detected.
in all the RNA samples obtained from laser microdissected cells. The use of gene-specific primers that span introns excluded the possibility of genomic DNA in total RNA samples used for RT-PCR analyses. The absence of an amplified product in RT-negative reactions also excluded any DNA contamination in RNA samples obtained from laser microdissected cells (results not shown). Finally, OsCPK4 transcripts were detected in RNA samples obtained from the two cell types captured from fungal-inoculated and control roots, this observation further supporting the integrity of the RNA samples used in this study (Additional file 2: Figure S2).

When comparing the results obtained on OsCPK18 expression in laser microdissected cells (Figure 2F) and whole roots (Figure 1), an apparent contradiction is observed. Thus, OsCPK18 transcripts were not detected in isolated cells from mock-inoculated roots (Figure 2F) whereas RT-qPCR analysis revealed OsCPK18 expression in whole roots (Figure 1A, upper panel). This finding could be explained taking into account the plant material and experimental approach used in these studies. In this work, only two cell types of the root were harvested for LMD-related analyses (epidermal and cortical cells). Thus, the detection of OsCPK18 expression in whole mock-inoculated roots could be due to the presence of cell types constitutively expressing OsCPK18 that were not analyzed with LMD (i.e. cells from the central cylinder). Additionally, transversal sections were routinely made at approx. 2 cm from the root tip. Thus the observed expression of the OsCPK18 gene in regions of the rice root other than that used for laser microdissection (i.e. meristems) might well account for the observed OsCPK18 expression in mock-inoculated whole roots. This observation also illustrates the fact that results obtained in gene expression by using whole roots might often be misinterpreted and spatial differences in gene expression might not be perceived by using whole roots. Clearly, a more detailed analysis of OsCPK18 expression during growth and development of the rice root is needed.

Subcellular localization of OsCPK18

Onion epidermal cells are widely used as a convenient system in which to evaluate the subcellular location of GFP-tagged proteins. Accordingly, the subcellular localization of OsCPK18 was investigated in onion epidermal cells transiently expressing gene fusions to the green fluorescent protein (GFP) (Figure 3A). Confocal microscopy of transformed onion cells revealed that OsCPK18-GFP localizes to the cell periphery, likely the plasma membrane (Figure 3B). As expected, onion cells expressing the GFP gene showed fluorescence distributed throughout the cell (Figure 3C).

Onion epidermal cells are also particularly useful for analysis of plasma membrane proteins because the environmental conditions can be manipulated to cause plasmolysis and partial separation of the plasma membrane from the cell wall. The onion epidermal cells were plasmolyzed after being transformed with OsCPK18-GFP. In plasmolyzed onion cells, the OsCPK18-GFP displayed a pattern consistent with its location in the plasma membrane of the shrunken protoplast (Figure 3D). Under these conditions, protoplast pull
away from the cell wall, leaving large numbers of thin plasma membrane bridges, known as Hechtian strands, firmly anchored to the cell wall (Figure 3E).

Analysis of the amino acid sequence of OsCPK18 shows that the OsCPK18 polypeptide possess a N-terminal myristoylation site at the Gly residue at position 2 (Gly2) suggestive of N-myristoylation. The need for this lipid modification to promote and stabilize membrane association of certain CPKs has been experimentally demonstrated [37]. To address the role of the myristoylation site of OsCPK18 in plasma membrane association, a mutation at the N-terminal myristoylation site (MGNTCVGPS) of the OsCPK18 polypeptide was made. The Gly2 was converted to Ala (G2A, referred to as OsCPK18G2A) and fused to GFP (Figure 3A). Transient expression in epidermal onion cells showed that the Gly2 mutation abolished the plasma membrane localization of OsCPK18 (Figure 3F). Instead, a distribution throughout the cell was observed for the mutated version of OsCPK18 similar to that of the GFP alone. These findings suggest that the N-terminal myristoylation site is required for subcellular localization of OsCPK18 at the plasma membrane.

Phylogenetic analysis of cpk genes
In this work, the evolutionary relationships among CPKs from rice and known CPKs from other plant species establishing association with AM fungi was determined. For this analysis, the full-length CPK protein sequences from cereal species, namely wheat and maize, as well as CPKs so far characterized in the model symbiotic species of Medicago were used. As previously mentioned, the rice genome contains 31 CPK genes which classify into four major phylogenetic groups (I-IV) [39,40]. Known CCaMK protein sequences from rice, wheat and Medicago were also considered. In this respect, the rice genome contains a single CCaMK gene [39]. As Arabidopsis is not a host for AM fungi, this species was not included in the phylogenetic analysis.

Phylogenetic trees of CPK and CCaMK proteins were constructed based on the neighbor-joining method (Figure 4) or the maximum parsimony method (Additional file 3: Figure S3). The alignment of the various proteins used for construction of the phylogenetic tree is presented in Additional file 4). Similar to what was previously reported [40], the rice CPKs clustered into four distinct phylogenetic groups (Figure 4). Four distinct CPKs, OsCPK18, OsCPK4, OsCPK30 and OsCPK31, cluster into an independent clade of CPKs, the Group IV, which appears to have diverged significantly from the other rice CPK sequences. Noticeably, results here presented show that OsCPK18 and OsCPK4 are both up-regulated by the AM fungus G. intraradices, these particular CPKs belonging to Group IV of rice CPKs. As for the other members of the Group IV of rice CPKs, no expression could be detected in the rice roots for OsCPK31, whereas OsCPK30 exhibited a low expression but no responsiveness to AM inoculation.

Some interesting observations came from the phylogenetic analysis of CPK and CCaMK proteins. Firstly, OsCPK18 and OsCPK4 appear to be closely related to the AM-associated MtCDPK1 (Figure 4). Secondly, Group IV of rice CPKs and CCaMKS are closely related each other. Indeed, Group IV of rice CPKs appears to be more related to CCaMKs than to the other rice CPKs. Here, it is worthwhile to mention that the essential function of MtCCaMK and OsCCaMK during the mycorrhizal symbiotic association is well documented [18,23]. Finally, the OsCPK18 is clearly related to TaCPK6, one of the 20 CPKs described in wheat [45].

Sequence analysis of the OsCPK18 and OsCPK4 promoters
Knowing that the OsCPK18 and OsCPK4 genes are transcriptionally activated in response to inoculation with
the AM fungus *G. intraradices*, it was of interest to investigate whether symbiosis-related cis-elements are present in the promoter region of these genes. The OsCPK18 and OsCPK4 promoter analysis was carried out using the PLACE algorithm [46] and extended to genes that are known to be required for both AM and rhizobial root nodule symbioses, such as the MtCPK1 and MtCCaMK genes from *M. truncatula* and the OsC-CaMK from rice.

Analysis of the 2 kb promoter region of the OsCPK18 and OsCPK4 genes revealed the presence of the CTCTT element (NODCON2GM) which is found up to five and six times in the OsCPK18 and OsCPK4 promoter, respectively (Figure 5 and Additional file 5: Tables S1 and S2). The NODCON2GM as well as the NODCON1GM element (AAAGAT) are characteristic motifs of promoters from genes that are regulated during root nodule and AM symbiosis. These motifs are also part of the "organ-specific element" (OSE) sequence [47]. The MtCPK1, OsCCaMK and MtCCaMK promoters contain several copies of the NODCON1GM and NODCON2GM consensus sequences.

Interestingly, multiple copies of the ABRE-related consensus motif [(C/A)ACG(T/C)G(T/G/C), ABRERATCAL] were present in the proximal region of the OsCPK18 promoter (Figure 5 and Additional file 5: Tables S1 and S2). The ABRE-related motif is a cis-element identified in the upstream region of 162 Ca²⁺-responsive up-regulated genes [48]. Furthermore, up to three copies of the CCGG-BOX element (GCGGCGGC) are found in the Oscpk18 promoter, this element being involved in Ca⁺⁺/calmodulin-regulated gene expression [49] (Figure 5 and Additional file 5: Tables S1 and S2). The OsCPK4 promoter region contains one copy of the ABRE-related motif element. The G(G/A/C/T)ATAT(G/A/C/T)C (P1BS element) was recognized in the OsCPK4, MtCPK1 and MtCCaMK promoters (Figure 5 and Additional file 5: Tables S1 and S2). This element is found in the upstream region of phosphate starvation responsive genes from several plant species [50].

Finally, the OsCPK18 and OsCPK4 promoters harbor multiple stress-related cis-acting elements, including elements that are known to confer responsiveness to pathogen-regulated genes. Some of them were represented many times in these promoters, such as the TGAC-containing W box of WRKY transcription factors (Additional file 5: Tables S1 and S2). In line with this, we recently reported the activation of defense-and stress-related genes during colonization of rice roots by *G. intraradices* [4]. Whether the expression of the OsCPK18 and OsCPK4 genes is regulated during pathogen infection in roots remains to be determined.

Overall, this study revealed the presence of symbiotic-related motifs, as well as putative elements related to Ca²⁺ regulation of gene expression, in the promoter region of the OsCPK18 and OsCPK4 genes. This observation is consistent with the observed induction for the two CPK genes in AM-inoculated rice roots.

**Discussion**

In this work, the expression of CPK genes was monitored during the early stages of the AM symbiosis in rice. The OsCPK18 and OsCPK4 consistently showed up-regulation in response to AM inoculation. Evidence is also presented on the transcriptional activation of OsCPK18 and OsCPK4 expression by diffusible molecules produced by *G. intraradices*. When comparing the expression profiles of the rice CPK and CCaMK genes, it appears that activation of the two CPK genes (OsCPK18 and OsCPK4) occurs earlier than that of OsCCaMK pointing to a role for these particular rice CPK genes at the early stages of the symbiotic process. The observation that the OsCPK18, OsCPK4, OsCCaMK, MtCPK1 and MtCCaMK genes share symbiotic-related cis-elements in their promoters is also indicative of the transcriptional regulation of these genes as part of the signaling mechanisms involved in the AM symbiosis in rice. An expanded view of OsCPK18 gene expression came from expression studies in laser microdissected
cells isolated from rice roots. At 4 days post-inoculation with *G. intraradices*, OsCPK18 was detected in cortical cells and not in epidermal cells.

Clearly, the specificity of a CPK functioning in a given signaling pathway may be achieved not only by a differential pattern of expression but also by targeting of the CPK protein to a particular subcellular compartment. Along with this, CPK proteins appear to be widely distributed among subcellular compartments including cytosol, peroxisome, plasma membrane, oil bodies and nucleus, as well as in association with actin filaments, mitochondria and the endoplasmic reticulum [33]. Our results in transformed onion cells clearly demonstrated that OsCPK18 localizes to the plasma membrane. Moreover, the association of OsCPK18 to the plasma membrane is possibly linked to N-terminal myristoylation of this protein.

Knowing that CPKs act as Ca$^{2+}$ sensors in plant signaling, and that Ca$^{2+}$ plays an important role in the AM symbiosis, a function of OsCPK18 as a Ca$^{2+}$ sensor during the AM-induced host responses to AM fungi can be envisaged. Thus, perception of the fungal-produced symbiotic signal(s) would activate downstream signaling events required for the establishment of the symbiotic association, including the cytoplasmic and nuclear Ca$^{2+}$ spiking responses [9-11]. Alterations in the Ca$^{2+}$ level would be itself a major factor in mediating up-regulation of OsCPK18 gene expression in the nucleus, as judged by the presence of the Ca$^{2+}$-responsive *cis*-elements in the OsCPK18 promoter region [48]. In line with this, previous studies in Arabidopsis revealed the presence of ABRE-related sequences in Ca$^{2+}$-responsive genes, and exclusively in up-regulated Ca$^{2+}$-responsive genes [48]. Tetramers of the ABRE-*cis* element are sufficient to confer this transcriptional activation in response to Ca$^{2+}$ transients. The presence of multiple Ca$^{2+}$-responsive *cis*-regulatory elements in the promoter region of the OsCPK18 gene (e.g. ABRE-related and CGCG-box elements) favors the possibility of a Ca$^{2+}$-mediated up-regulation of OsCPK18 gene expression. The identity of the transcription factors that respond to rapid transient Ca$^{2+}$ signals and that subsequently activate gene expression through ABRE-related *cis*-elements remains to be determined.

In addition to its transcriptional activation, a direct regulation of the OsCPK18 enzyme activity by Ca$^{2+}$ can be expected. Thus, it is well known that the activity of CPKs is regulated by the binding of calcium to its intrinsic calmodulin-like domain. At basal Ca$^{2+}$ concentrations, the functional autoregulatory domain acts as a pseudosubstrate that inhibits the kinase activity of CPKs (autoinhibited structure). In response to transient increases in the level of cellular Ca$^{2+}$, CPKs undergo conformational changes that activate their kinase activity (calcium-bound structure) [51]. It is then reasonable to assume that the plasma membrane-localized OsCPK18 protein sense the AM-induced increase in cytoplasmic Ca$^{2+}$ levels and transduce this signal into phosphorylation processes. The OsCPK18-mediated signaling processes might then be crucial for root colonization and accommodation of the fungal symbiont in the root cortex. The identification of downstream targets of the OsCPK18 kinase activity requires, however, further investigation.

On the other hand, our phylogenetic analysis of CPKs and CCaMK of plant species that are able to establish mycorrhizal associations revealed that Group IV of CPKs and CCaMK are closely related each other pointing to an evolutionary relationship between the two families of protein kinases. In other studies carried out in the green alga it was proposed that CCaMK originated through gene duplication from CPK during green alga evolution [52]. Altogether, these findings are in clear support a functional specialization of members of the Group IV of CPKs and their relatedness with CCaMK functioning. Adaptation steps probably occurred in different plant species that determined their functional specialization and symbiosis-specific regulation.

The current work also provides a foundation for further functional investigation of the complex CPK family in relationship to the mycorrhization ability in another cereal species, such as wheat. Thus, the phylogenetic analysis of CPKs revealed that OsCPK18 and OsCPK4 are closely related to the wheat TaCPK6 protein as well as to the *Medicago* MtCDPK1 protein. For MtCDPK1 a role during the establishment of the AM symbiosis is well documented [53]. It is then tempting to speculate that the TaCPK6 gene might exhibit an AM-regulated expression pattern in wheat plants.

An intriguing aspect is the presence of three Arabidopsis proteins in Group IV of CPKs [39], even though Arabidopsis is not a host for AM fungi. To this point, it has been proposed that genes required for other aspects of plant development might have been recruited to function in symbiotic pathways. In line with this, inactivation of the *MtCDPK1* gene is associated to a significant reduction of rhizobial and mycorrhizal symbiosis and also results in stunted roots and short root hairs in *M. truncatula* [53]. In other studies, impairment of root hair development results in defective symbiotic interactions in *L. japonicus* [54]. Then, the Arabidopsis CPKs within Group IV of CPKs might play a role in normal processes during root growth and development. The finding of SYM genes in species that do not associate with AM fungi (e.g. Arabidopsis and *Physcomitrella*), also supports that specific genes functioning in normal developmental processes in roots might also regulate mycorrhizal infection. If so, this fact, would explain the
observed expression of OsCPK18 in experiments carried out on whole roots by RT-qPCR.

Conclusions
This study provides a new view of the molecular mechanisms involved in the AM symbiosis in rice while defining an OsCPK18-mediated signaling pathway functioning during this process. The rapid activation of OsCPK18 expression in response to AM inoculation, its expression being also induced by fungal-secreted signals, together with the observed plasma membrane localization of OsCPK18, suggest that OsCPK18 might play a role in perception and/or recognition of the AM fungus in rice. Compared to legume species, less effort has been invested in the characterization of the AM symbiotic interaction in this important crop species. OsCPK18 might be considered as a marker for the presymbiotic phase of the symbiotic process that might play a preferential role in the root cortex. The identification of additional components of the AM-induced signaling processes in which OsCPK18 participates can be now approached. A major challenge for the future research is to determine whether interconnections and synergistic functions exist between CPKs and SYM components, this interplay determining recognition and compatibility between the two symbiotic partners.

Methods
Plant material and growth conditions
Rice (Oryza sativa cv Nipponbare) was used as the experimental material. Seeds were surface sterilized with 70% ethanol for 1 min, sodium hypochlorite (30% v/v) for 30 min, and extensively washed with sterile water (four times, 10 min each). Seeds were germinated in agar (0.4%) prepared with minimal medium. Since Millipore membranes are permeable to diffusible molecules, the root cells can perceive fungal signals in the double sandwich method even though physical contact between the two symbionts does not occur. Control seedlings were inoculated with sterile water.

Tissue preparation and laser microdissection
The method previously described [43] was adapted for the isolation of cells from G. intraradices-inoculated and mock-inoculated rice roots. Root pieces of 4 - 8 mm in length were dissected with a razor blade and immediately transferred into freshly prepared Methacarn solution (absolute methanol/chloroform/glacial acetic acid 6:3:1). Roots were maintained in the fixative solution overnight at 4°C, and subsequently dehydrated in a graded series of ethanol at 4°C: 50, 70 and 90% in sterile water and 100% ethanol, followed by isopropanol (twice), with each step on ice for 1 h. The isopropanol was replaced gradually with paraffin (Paraplast Plus; Sigma Aldrich, St. Louis). Transverse root sections of 10-15 μm were made using a Reichert Jung 2050 SuperCut Motorized Microtome (Leica, Arnsberg, Germany). Ribbons were arranged on RNase-free, UV-treated, PEN-membrane 2.0 μm slides. Slides were kept in a slide warmer at 40°C until dry and stored at 4°C and used within two days.

The Leica LD6000 Laser Microdissection system (Leica, Bannockburn, IL, USA) was used for laser microdissection (LMD). Just before use, the paraffin sections were deparaffinised in a neoclear (Merck, Darmstadt, Germany) treatment for 10 min followed by 100% ethanol for 2 min, and then air dried. The deparaffinised slides were placed face down on the microscope. The tissues were visualized on a computer monitor through a video camera. Epidermal and cortical cells were marked and then cut using a UV laser (337-nm wavelength). Target cells were collected without any extra forces into the cap of a microcentrifuge (RNase-free PCR tube caps). For each cell type, we isolated at least 1500 cell sections per biological replicate, and two independent biological replicates were made. After collection, 50 μl of RNA extraction buffer from the PicoPure kit (Arcturus, Sunnyvale, CA, U.S.A.) were added. Samples were incubated at 42°C for 30 min, centrifuged at 800 g for 2 min, and stored at -80°C until RNA isolation.
RNA isolation

Total RNA was extracted from whole roots at different times after inoculation with *G. intraradices* spores, as well as from mock-inoculated roots, using the TRI-ZOL® Reagent (Invitrogen, Carlsbad, CA, USA). For each time point, roots from at least 12 individual plants were collected. Three independent experiments were carried out. The first cDNA was synthesized from DNase-treated total RNA (1 μg) with M-MLV (Moloney-Murine Leukemia Virus) Reverse Transcriptase (Invitrogen, Carlsbad, CA, USA). Aliquots of the resulting RT reaction product were used as template for PCR analysis.

Leukemia Virus) Reverse Transcriptase (Invitrogen, Carlsbad, CA, USA). Essentially, the DNase-treatment was not performed on the kit column but the treatment was performed using a modified PicoPure kit protocol (Arcturus, Sunnyvale, CA, USA). The aliquots of the resulting RT reaction product were used as template for PCR analysis.

RNA isolations from laser microdissected cells were performed using a modified PicoPure kit protocol (Arcturus, Sunnyvale, CA, USA). The aliquots of the resulting RT reaction product were used as template for PCR analysis.

Phylogenetic analysis

Quantitative real time PCR (RT-qPCR) analyses were carried out in optical 96-well plates in a LightCycler® 480 Real-Time PCR System (Roche) according to the following program: 10 min at 95°C, followed by 45 cycles of 95°C for 10 s, 60°C for 30 s, and an additional cycle of dissociation curves to ensure an unique amplification. The reaction mixture contained 10 μl 2× SYBR Green Master mix reagent (Roche, Mannheim, Germany), 2 μl cDNA sample, and 300 nM of each gene-specific primers, in a final volume of 20 μl. Primers used for RT-qPCR are indicated in Additional file 6: Table S3. Details on RT-qPCR analysis following the MIQE guidelines [55] are included in Additional file 7. Routinely, three replicate reactions were used for each sample. Data were normalized with OsAct1 as internal control. The average CT values from triplicate PCRs were normalized to the average CT values for the OsAct1 gene from the same RNA preparations. Three independent biological replicates were analyzed. For each biological material, three technical replicates were made for RT-qPCR analysis.

In this work, semi-quantitative reverse-transcription-polymerase chain reaction (RT-PCR) was carried out to investigate the expression pattern of the rice *CPK* genes, as well as the rice *SYM* genes, namely the *OsSYM1*, *OsSYM2*, *OsSYM3*, *OsSYM4*, *OsSYM5*, *OsSYM6*, *OsSYM7*, and *OsSYM8* genes. (Additional file 6: Table S4 and Additional file 9: Supplementary Methods).

For RT-PCR analysis of RNA samples obtained from laser microdissected cells, a one-step RT-PCR was conducted according to the manufacturer’s instructions (Qiagen Gmbh, Hilden, Germany). A 20 μl reaction was prepared, containing the following reagents for each reaction: 4 μl of (5X) Qiagen one-step RT-PCR buffer, 0.5 μl of Qiagen one-step RT-PCR enzyme mix, 0.5 mM of each dNTP, 0.25 μM of each primer, 21 ng of total RNA and RNase free water to 20 μl.

Promoter analysis

Sequences 2000 bp upstream of the selected *cpk*’s genes were retrieved from NCBI database. Known plant motifs were obtained from the PLACE database http://www.dna.affrc.go.jp/PLACE/.

Biological cell transformation and imaging by confocal microscopy

To investigate the subcellular localization of OsCPK18, the green fluorescent protein (GFP) gene was translationally fused to the C-terminal end of the OsCPK18 sequence. For this, a GFP gene lacking the N-terminal signal peptide and C-terminal HDEL sequences was initially obtained by PCR from the *m-GFP5-ER* sequence.
OsCPK18 DNA fragment to be translationally fused to the GFP gene was generated by PCR amplification. A BglIII site was introduced at the 3’ end of the PCR-amplified Osckpt18 DNA fragment. Equally, a SpeI site was introduced at the 5’ end of the Osckpt18 sequence. The full-length OsCPK18 cDNA sequence was PCR-amplified from clone J023148F12 obtained from the KOME (Knowledge-based Oryza Molecular biological Encyclopedia) database using the following primers: 5’-ATACGTAGATGGAACCTGCTGCTCTCC-3’ (forward) and 5’-ATAGATCTACCTGGTGTTGGCGATCTGTGAACACTCCT-5’ (reverse) (underlined sequences indicate the SpeI and BglIII restriction sites; bold fonts indicate the residual tail of one glycine and three prolines to ensure the fusion). To obtain the chimeric OsCPK18-GFP gene, the PCR-amplified OsCPK18 DNA fragment was digested with SpeI and BglIII and cloned into the XbaI/BamHI-digested pP35S:mGFP:nos plasmid resulting in plasmid pP35S:OsCPK18-GFP:nos.

A chimeric gene in which a mutated version of OsCPK18 gene was fused to the GFP gene was also prepared. For preparation of the mutated version of the OsCPK18 gene in which the Gly residue at position 2 was converted to Ala (G2A), the pP35S:OsCPK18-GFP:nos plasmid was used as the template for PCR. For this, a forward primer with a single change (bold font) 5’-ATACGTAGATGGAACCTGCTGCTCTCC-3’ (forward) and 5’-ATACGTAGATGGAACCTGCTGCTCTCC-3’ (reverse) (underlined sequences indicate the SpeI and BglIII restriction sites; bold fonts indicate the residual tail of one glycine and three prolines to ensure the fusion). To obtain the chimeric OsCPK18-GFP gene, the PCR-amplified OsCPK18 DNA fragment was digested with SpeI and BglIII and cloned into the XbaI/BamHI-digested pP35S:mGFP:nos plasmid resulting in plasmid pP35S:OsCPK18-GFP:nos.

Additional material
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Authors’ contributions
LCS carried out most of the experimental work and data analyses. JGA participated in laser microdissection and gene expression studies. BSS coordinated the design and execution of this study and wrote the manuscript. PB also participated in the design of this study and critically revised the manuscript for important intellectual content. All authors read and approved the final manuscript.

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