

# A PAS-Containing Histidine Kinase is Required for Conidiation, Appressorium Formation, and Disease Development in the Rice Blast Fungus, *Magnaporthe oryzae*

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## ABSTRACT

Rice blast disease, caused by the ascomycete fungus *Magnaporthe oryzae*, is one of the most important diseases in rice production. PAS (period circadian protein, aryl hydrocarbon receptor nuclear translocator protein, single-minded protein) domains are known to be involved in signal transduction pathways, but their functional roles have not been well studied in fungi. In this study, targeted gene deletion was carried out to investigate the functional roles of the PAS-containing gene *MoPAS1* (MGG\_02665) in *M. oryzae*. The deletion mutant  $\Delta MoPAS1$  exhibited easily wettable mycelia, reduced conidiation, and defects in appressorium formation and disease development compared to the wild type and complemented transformant. Exogenous cAMP restored appressorium formation in  $\Delta MoPAS1$ , but the shape of the restored appressorium was irregular, indicating that *MoPAS1* is involved in sensing the hydrophobic surface. To examine the expression and localization of *MoPAS1* in *M. oryzae* during appressorium development and plant infection, we constructed a *MoPAS1*:GFP fusion construct. *MoPAS1*:GFP was observed in conidia and germ tubes at 0 and 2 h post-infection (hpi) on hydrophobic cover slips. By 8 hpi, most of the GFP signal was observed in the appressoria. During invasive growth in host cells, *MoPAS1*:GFP was found to be fully expressed in not only the appressoria but also invasive hyphae, suggesting that *MoPAS1* may contribute to disease development in host cells. These results expand our knowledge of the roles of PAS-containing regulatory genes in the plant-pathogenic fungus *M. oryzae*.

## ARTICLE HISTORY

Received 27 August 2019  
Revised 29 October 2019  
Accepted 31 October 2019

## KEYWORDS

*Magnaporthe oryzae*; rice blast; PAS domain

## 1. Introduction

Fungi have evolved to sense ambient environmental signals, such as air, light, and host surface features, for appropriate development and survival [1]. For example, G-protein-coupled receptors (GPCRs), one of the largest families of transmembrane receptors, are involved in sensing environmental signals and activating signal transduction pathways [2]. The PAS (period circadian protein, aryl hydrocarbon receptor nuclear translocator protein, single-minded protein) domain is involved in sensing signals, and is present in many signaling proteins, such as histidines (HKs) and serine/threonine (S/T) kinases, as well as voltage-gated ion channels [3,4]. The PAS domain was first discovered in *Drosophila* proteins, and has since been found in organisms ranging from bacteria to humans [3,5]. In bacteria, PAS domains are known to associate with histidine kinase (HK) sensor proteins [6]. HKs were initially believed to exist only in bacteria, but in the 1990s it was revealed that HKs also regulate essential

processes in fungi [7]. In addition to their sensory role, PAS domains also modulate protein–protein interactions in response to stimuli, which allows for complex cellular signaling networks [8].

PAS domains are found in previously characterized regulators in some fungi. For example, PAS domains are present in the serine/threonine protein kinases PSK1 and PSK2 in yeast [9]. In yeast, PSK1 regulates superoxide dismutase-1 (SOD1) to protect against oxidative stress, and PSK2 acts as a nutrient-sensing kinase [10]. In the filamentous fungus *Neurospora crassa*, DCC-1, a putative HK, is involved in the regulation of conidiation, perithecial development, and carotenogenesis [11]. In the ascomycete fungus *Aspergillus nidulans*, phytochrome FphA regulates sexual development and mycotoxin formation [12]. These studies suggest important roles for PAS-containing proteins in diverse aspects of fungal development.

*Magnaporthe oryzae* is a filamentous plant pathogen that causes rice blast disease. This fungus

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has been used as a model organism for studying the molecular basis for fungal development and pathogenicity [13]. On rice leaves, conidia germinate and develop specialized infection structures called appressoria in response to host signaling factors. Sensing host signaling factors and activating signaling pathways are known to be necessary for *M. oryzae* to infect rice. cAMP-dependent protein kinase signaling is needed for appressorium formation in response to plant signaling factors in *M. oryzae* [14,15]. Pth11, a GPCR, is also known to be important in hydrophobic surface sensing for appressorium formation [16,17]. Recently, HKs were functionally characterized in *M. oryzae* [18]. In this previous study, the deletion mutant  $\Delta$ *Mohik5* was unable to form appressoria and was therefore completely nonpathogenic.

Previously, we identified conidiation-related genes from microarray analysis, among which attention has been paid to *MoPAS1* (MGG\_02665) due to potential important roles in fungal development and pathogenicity [19]. We generated a targeted gene deletion mutant for *MoPAS1*, named as  $\Delta$ *Mopas1*. Deletion of *MoPAS1* resulted in pleiotropic defects in conidiation, surface hydrophobicity of mycelia, appressorium development, and disease development. The defect of  $\Delta$ *Mopas1* in appressoria formation on hydrophobic coverslips was restored by cAMP treatment, which suggested that *MoPAS1* plays an important role in surface sensing. Collectively, the results showed that *MoPAS1* plays critical roles in preinfection-associated developmental stages and disease development in *M. oryzae*.

## 2. Materials and methods

### 2.1. Phylogenetic analysis

All sequence information was obtained from the Magnaporthe genome database of the Broad Institute (<http://www.broadinstitute.org>), the Comparative Fungal Genomics Platform (CFGP) online database (<http://cfgp.snu.ac.kr>), and the BLAST program provided by the National Center for Biotechnology Information (NCBI, <http://www.ncbi.nlm.nih.gov>). Sequence alignment was performed using ClustalW program in MEGA 6.0 [20,21]. Phylogenetic analysis was performed using the neighbor-joining method (1000 bootstrap replicates). Domain structure analysis was performed using InterPro Scan v53.0 (<http://www.ebi.ac.uk/interpro/>) [22]. Domain structure analysis was performed using InterPro Scan (<http://www.ebi.ac.uk/interpro/>) [22]. The primers used in this study (Table 1) were designed using the PrimerQuest Design Tool (<http://sg.idtdna.com/site>), and purchased from Bioneer (Daejeon, Korea).

### 2.2. Fungal strains, culture conditions, and phenotypic characterization

The *M. oryzae* wild-type KJ201 and transformants used in this study were cultured on oatmeal agar (OMA, 50 g of oatmeal and 25 g of agar per liter) or V8 juice agar (V8A, 80 mL of V8 juice, 310  $\mu$ L of 10 N NaOH solution, and 15 g of agar per liter) at 25 °C under continuous light. Conidiation of *M. oryzae* was measured by counting the number of conidia harvested from 7-d-old OMA plates using a hemocytometer. For microscopic observation of conidiation, mycelial agar plugs from 4-d-old OMA plates were transferred on slide glasses using a cork borer and covered with cover slips. After 1 d, conidiation was observed under a light microscope. To measure conidial germination and appressorium formation, we collected conidia from 7-d-old OMA plates and placed drops (20  $\mu$ L) of conidial suspensions ( $5 \times 10^4$  conidia/mL) on hydrophobic cover slips and incubated them in a moistened box. Germination was examined by counting the number of germinated conidia from 100 conidia per coverslip after 2 h and appressorium formation was measured after 16 h. For cAMP assays, we prepared cAMP solution (10 mM) in sterile distilled water (SDW), and cAMP solutions were added to the conidial suspensions to a final concentration of 5 mM after a 2-h incubation. For surface hydrophobicity assays, drops (10  $\mu$ L) of SDW or 0.2% SDS were placed on wild-type and mutant strains grown on OMA plates for 10 d. Experiments were repeated three times with three replicates for each repeat. All data were processed using the SigmaStar statistical software package (SPSS Science, Chicago, IL).

### 2.3. Fungal transformation for gene knock-out and complementation

Approximately 1.0-kb flanking regions of *MoPAS1* were amplified using primers MoPAS1\_5F/5R and MoPAS1\_3F/3R from wild-type genomic DNA. The hygromycin phosphotransferase gene (*HPH*) cassette was amplified from the plasmid pBCATPH using HPH\_F/R primers. The amplified fragments were fused by double-joint PCR [23]. The fused construct was amplified using MoPAS\_NF/NR primers. The amplified constructs were transformed to wild-type protoplasts using the polyethylene glycol (PEG)-mediated transformation method [24]. Hygromycin-resistant transformants were screened on transformation agar (TB3, 200 g of sucrose, 3 g of yeast extract, 3 g of casamino acids, 10 g of glucose, and 8 g of agar per liter) containing Hygromycin B (200  $\mu$ g/mL). Targeted gene deletion was confirmed using Southern blot hybridization and RT-PCR [25,26]. For complementation,

**Table 1.** Primers used in this study.

Name	Primer sequence 5' to 3'
MoPAS1_5F	CTCACGTACCAACCTGCAAA
MoPAS1_5R	CCTCCACTAGTCCAGCCAAGCCCCACAACCCATCGGTCTTGATAG
MoPAS1_3F	GCCGGATTATCGTCTTCTTTGCGCGATCATCGTCTCTTTT
MoPAS1_3R	ACGACCTTGTGACTGGTACT
MoPAS1_NF	GGTCCCAATTGATGCCAAAG
MoPAS1_NR	TGCTGATACGAACACCAGATAG
MoPAS1_SF	GTCTACTGGACCCGAACACTA
MoPAS1_SR	GCCAGAAAGAAGGACGATGAATCCGG
MoPAS1_PF	CCCTACGCATGGATGGATT
MoPAS1_PR	GCAGGGTAGTGATAACGAGAAC
MoPAS1_cmF	CCATGGATGTCAGCAGATTTTGCCA
MoPAS1_cmR	GGATCCTCACTCGCTGCCGGCAGGT
MoPAS1_RTF	CTCGAGGGTAAAGCAAGTTCTC
MoPAS1_RTR	CGGCCACGAAGTCTGAATATAG
HPF_F	GGCTTGGCTGGAGCTAGTGAGG
HPF_R	CTCCGGAGCTGACATCGACACCAAC
Gen_F	AGAAGATGATATTGAAGG
Gen_R	CTTAAACAAGTGTACCTGTGC
Tub_F	TCGACAGCAATGGAGTTTAC
Tub_R	AGCACCAGACTGACCGAAGAC
MPG1_F	TCACCATCAACCAGCTCATC
MPG1_R	GGATGTTGACCAGACCAATCT
MHP1_F	GTCGATGCCGACAACCTCT
MHP1_R	GGGTGTTGCAGAGGATACC
GFP_F	CCACATCACTGCCAAGACAA
GFP_R	CTCGCTGCCGGCAGGTGAAGT
pIGPAPA_F	CCTGCCGGCAGCGAGATGGTGAGCAAGGGCGAGGA
pIGPAPA_R	TTGGCAGTGATGTGGCAACATACGAGCCGGAAGCA

*MoPAS1* with native promoters was amplified from wild-type genomic DNA using MoPAS1\_cmF/R primers and co-transformed to  $\Delta Mopas1$  protoplasts with plasmid pII99, which contains a geneticin-resistance cassette [27].

#### 2.4. RNA isolation, RT-PCR, and gene expression analysis

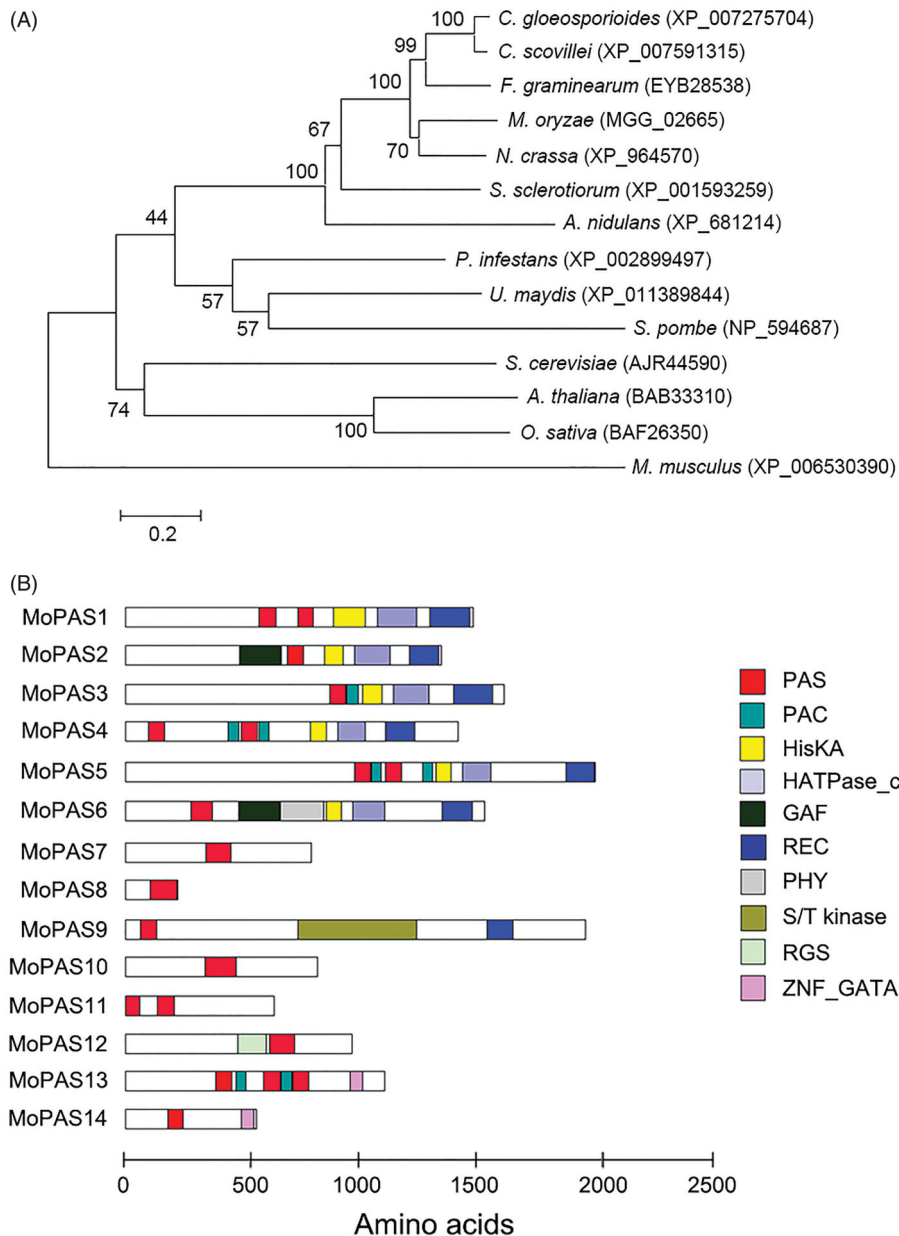
Total RNA was extracted from frozen fungal tissues using the Easy-Spin Total RNA Extraction Kit (Intron Biotechnology, Seongnam, Korea) according to the manufacturer's instructions. For reverse transcription PCR (RT-PCR), 5  $\mu$ g of total RNA was reverse transcribed into first-strand cDNA using oligo (dT) primers with the SuperScript III First-Strand Synthesis System Kit (Invitrogen Life Technologies, Carlsbad, CA). RT-PCR was performed in 20- $\mu$ L reaction mixtures using *Pfu* polymerase (Elpis, Daejeon, Korea). Cycling conditions were as follows: 3 min at 95 °C (1 cycle) followed by 20 s at 95 °C, 20 s at 57 °C, and 30 s at 72 °C (30 cycles). The  $\beta$ -tubulin (MGG\_00604) gene was used as a control. Quantitative real-time PCR was performed on the StepOne Real-Time PCR System (Applied Biosystems, Foster City, CA) using HIPI Real-Time PCR 2 $\times$  Master Mix (SYBR Green) (Elpis), and the relative abundance of *MPG1* and *MHP1* transcripts was measured as described previously [25]. The experiment was performed with three independent pools of tissues in two sets of experimental replicates.

#### 2.5. Pathogenicity assays

For spray assays, 3-week-old rice seedlings (cv. Nakdongbyeo) were sprayed with conidial suspensions ( $5 \times 10^5$  per mL) and incubated in a dew chamber 25 °C for 7 d [28]. For mycelial infection assays, mycelia agar plugs were placed on 4-week-old rice leaves cut into 8-cm lengths and incubated in a moistened box at room temperature for 7 d. To determine the penetration and infection ability of conidia, conidial suspensions ( $1 \times 10^5$  conidia/mL) were inserted into 4-week-old rice sheath tissue or placed on onion epidermis and incubated in a moistened box. Microscopic observation was performed at 24 and 48 h post-infection (hpi). Experiments were repeated three times with three replicates for each repeat.

#### 2.6. Localization of MoPAS1:GFP fusion protein

Overlap cloning was performed to generate the MoPAS1:GFP vector (Han *et al.*, 2015). A 4.4-kb fragment including a 2.4-kb ORF and 2.0-kb native promoter of *MoPAS1* was amplified using GFP\_F/R primers (Table 1). A 5.1-kb fragment from the plasmid pIGPAPA was amplified using pIGPAPA\_F/R primers. The amplified 2.0-kb and 5.1-kb fragments were cloned using the Overlap DNA Cloning Kit (Elpis Biotech, Korea), and the fused vector was transformed into wild-type [25]. Fluorescence microscopy images of MoPAS1:GFP localization were acquired using a Carl Zeiss Axio Image A2 microscope (Carl Zeiss Microscope Division, Oberkochen, Germany).



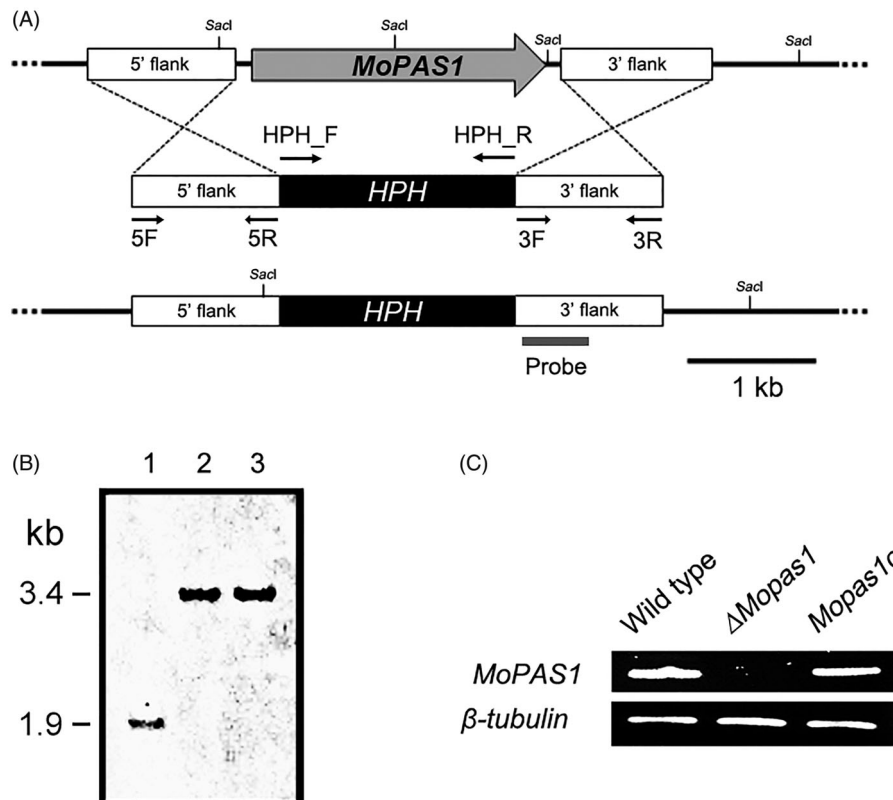
**Figure 1.** Phylogenetic analysis of MoPAS1. (A) Phylogenetic tree analysis of MoPAS1. A neighbor-joining tree of the amino acid sequence of PAS proteins. Numbers at nodes represent bootstrap values calculated from 1000 replicates. The scale bar indicates the number of amino acid differences per site. Sequence information was taken from the NCBI protein database. GenBank accession number for each protein is followed by species; (B) Predicted domain structure of MoPAS proteins. Sequence information was taken from the Magnaporthe genome database of the Broad Institute, and domain structure analysis was performed using InterPro Scan (<http://www.ebi.ac.uk/interpro/>). PAS (period circadian protein, aryl hydrocarbon receptor nuclear translocator protein, single-minded protein; IPR0000014), PAC (C-terminal to PAS motifs; IPR001610), HisKA (signal transduction histidine kinase; IPR003661), HATPase\_C (histidine kinase-like ATPase; IPR003594), GAF (domain present phytochrome and cGMP-specific phosphodiesterases; IPR003018), REC (signal receiver domain; IPR001789), PHY (phytochrome; IPR001294), S/T kinase (serine/threonine kinase; IPR008271), RGS (regulator of G protein signaling; IPR016137), ZNF\_GATA (zinc finger, GATA-type; IPR000679).

### 3. Results

#### 3.1. Phylogenetic analysis of MoPAS1

The *MoPAS1* (MGG\_02665) was predicted to encode a 1481 amino acids. To analyze genetic relationship of MoPAS1 with other organisms, a phylogenetic tree was constructed based on alignments of MoPAS1 amino acid sequences and related proteins in other organisms (Figure 1(A)). In the

phylogenetic tree, MoPAS1 was closely related to proteins in the subphylum Pezizomycotina filamentous fungi *Colletotrichum gloeosporioides*, *C. scovillei*, *Fusarium graminearum*, *M. oryzae*, *Neurospora crassa*, *Sclerotinia sclerotiorum*, and *Aspergillus nidulans* but distantly related to proteins in the subphylum Saccharomycotina or phylum Basidiomycota, which suggests that PAS1 proteins have evolved in a lineage-specific manner. Next, we searched for PAS



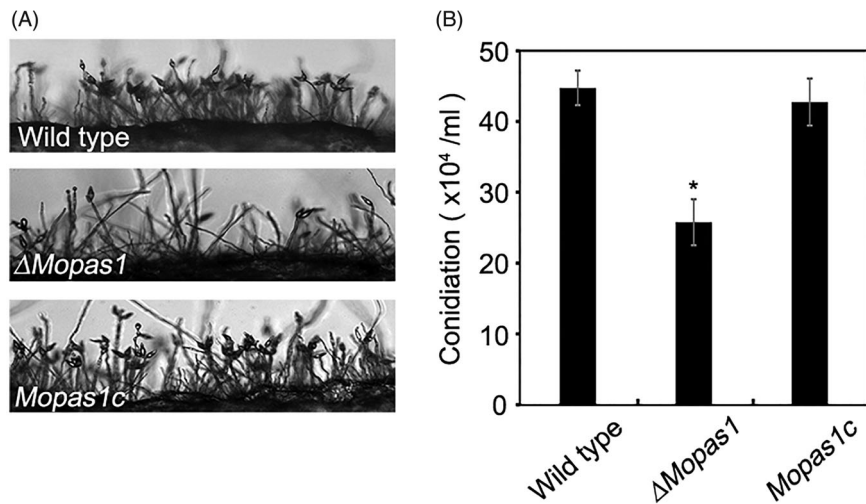
**Figure 2.** Gene deletion construct and identification of deletion mutants. (A) *MoPAS1* was deleted through the targeted gene replacement method; (B) Confirmation of *MoPAS1* deletion using Southern blot analysis. DNA samples were digested with *SacI*. A 1.9-kb band from wild-type KJ201 and a 3.4-kb band from knock-out mutants were produced. Lane 1, wild-type; lane 2–3, knock-out mutants; (C) Confirmation of *MoPAS1* deletion and complemented transformant *Mopas1c* using RT-PCR.

domains in the fungal database (fungiDB.org) and found several proteins containing PAS domains. Fourteen proteins containing PAS domains were found in *N. crassa* and *Fusarium graminearum*, whereas fewer were found in *Cryptococcus neoformans* (nine), *Aspergillus fumigatus* (seven), *Schizosaccharomyces pombe* (six), and *Saccharomyces cerevisiae* (three) (data not shown). A total of 14 proteins were found to have PAS domains in *M. oryzae* strain KJ201 (Figure 1). These proteins were named MoPAS1–MoPAS22. Each protein had one to three copies of the PAS domain, and proteins ranged from 225 to 1994 amino acids in length. In MoPAS1, there were two PAS domains, a HisKA (HK phosphoacceptor domain; IPR003661) domain, a HATPase\_C (HK-like ATPase; IPR003594) domain, and a REC (signal receiver domain; IPR001789) domain. Among the 14 proteins containing PAS domains from *M. oryzae* strain KJ201, PAC (C-terminal to PAS motifs; IPR001610), HisKA, HATPase\_C, GAF (domain present in phytochrome and cGMP-specific phosphodiesterases; IPR003018), REC, PHY (phytochrome; IPR001294), S/T kinase (serine/threonine kinase; IPR008271), RGS (regulator of G protein signaling; IPR016137) and ZNF\_GATA (zinc finger,

GATA-type; IPR000679) domains were present (Figure 1). In summary, PAS domains are present in the regulatory proteins of *M. oryzae* strain KJ201.

### 3.2. Targeted deletion of MoPAS1

To investigate the functional roles of *MoPAS1*, we generated a gene deletion mutant,  $\Delta Mopas1$  (Figure 2). The deletion construct was amplified by double-joint PCR. The construct is illustrated in Figure 2(A). The construct was used to transform protoplasts of *M. oryzae* strain KJ201 by a PEG-mediated transformation method. More than 100 putative deletion transformants were grown on TB3 agar containing 200  $\mu\text{g}/\text{mL}$  hygromycin B, and two candidates were selected by PCR screening using MoPAS1\_SF/SR primers (Table 1). Finally,  $\Delta Mopas1$  was selected based on Southern blot assay results. As illustrated in Figure 2(A), genomic DNA of the transformants was digested using *SacI* and hybridized with the probe. Southern blot analysis revealed that a 1.9-kb band was detected in the wild-type and a 3.4-kb band was detected in knock-out mutants (Figure 2(B)). Next, we confirmed the expression of *MoPAS1* by RT-PCR (Figure 2(C)). The expression

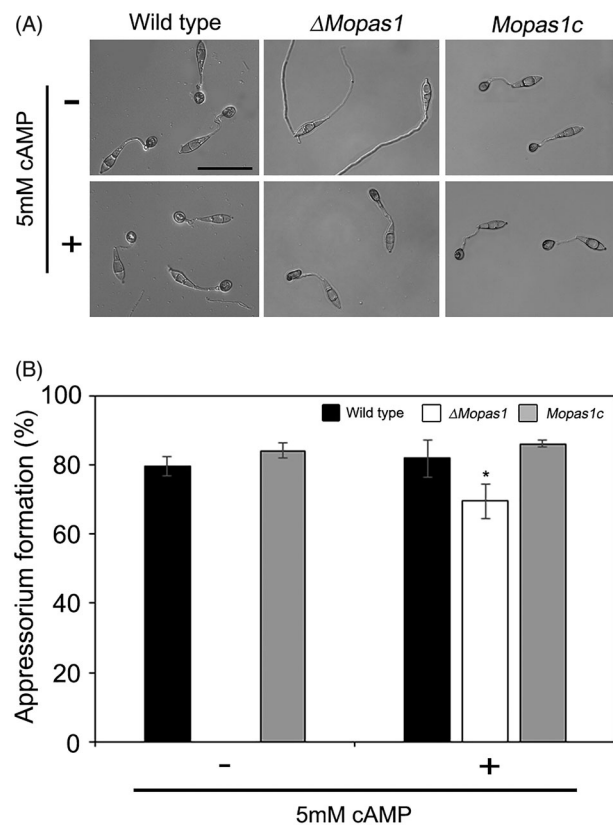


**Figure 3.** Conidiation and conidiophore differentiation. (A) Microscopic visualization of conidia and conidiophores in the wild-type strain,  $\Delta Mopas1$ , and  $Mopas1c$ ; (B) Quantitative measurement of conidia. Each strain was grown on OMA for 7 d, and conidia were counted using a hemocytometer. Asterisks represent significant differences from the wild-type strain (Tukey's test,  $p < .05$ ).

of *MoPAS1* was completely abolished in  $\Delta Mopas1$ , and was restored in  $Mopas1c$  (Figure 2(C)).

### 3.3. *MoPAS1* is essential for conidiation and appressorium development

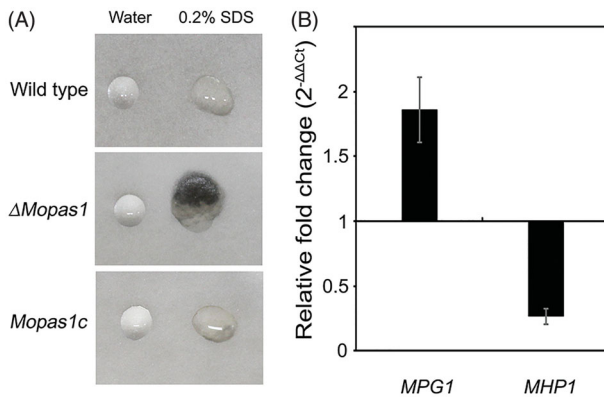
Conidia are important for disease dissemination in *M. oryzae*. To investigate the functional role of *MoPAS1* in conidiation, we assessed the conidiation of  $\Delta Mopas1$  on OMA media.  $\Delta Mopas1$  exhibited significantly reduced conidiation on OMA media (Figure 3(A,B)). At 7 d after inoculation on OMA, the wild-type strain produced approximately  $45 \times 10^4$  conidia/mL, whereas  $\Delta Mopas1$  only produced  $26 \times 10^4$  conidia/mL. This result means that *MoPAS1* is involved in conidiation. To investigate whether conidia of  $\Delta Mopas1$  are normal in appressorium formation, we incubated 20  $\mu$ L of conidial drops ( $5 \times 10^4$  conidia/mL) on hydrophobic coverslips. The conidia of wild-type germinated and formed melanized appressoria.  $\Delta Mopas1$  conidia also germinated on hydrophobic coverslips like wild-type but was unable to develop appressoria, which indicates that *MoPAS1* is involved in appressorium formation (Figure 4(A,B)). We then investigated whether the defect of  $\Delta Mopas1$  in appressorium formation is related to cAMP-dependent signal. We added exogenous cAMP solutions on the conidial drops on hydrophobic coverslips. The addition of exogenous cAMP restored appressorium formation in  $\Delta Mopas1$ . However, the restored appressoria of  $\Delta Mopas1$  were abnormally shaped, indicating that *MoPAS1* is involved in surface recognition and the morphogenesis of infection structures.



**Figure 4.** Appressorium formation. (A and B) Conidial suspensions of the wild-type strain,  $\Delta Mopas1$ , and  $Mopas1c$  were incubated for 12 hpi on hydrophobic surfaces. For cAMP assays, conidial suspensions of each strain were incubated for 12 hpi on hydrophobic surfaces. cAMP solutions were added to the conidial suspensions to a final concentration of 5 mM at 2 hpi. Asterisks represent significant differences from the wild-type strain (Tukey's test,  $p < .05$ ). Bar = 50  $\mu$ m.

### 3.4. *MoPAS1* is indispensable for surface hydrophobicity

Hydrophobins are low-molecular-weight proteins known to confer hydrophobicity in fungal conidia

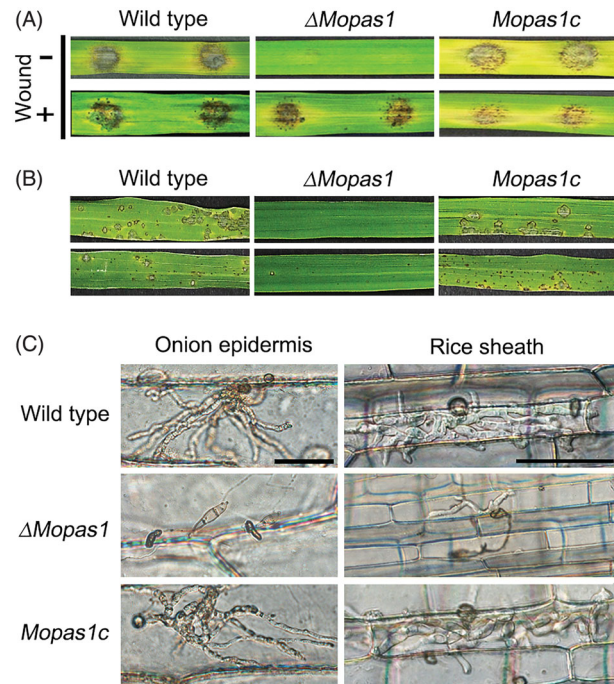


**Figure 5.** Surface hydrophobicity assessment of mycelia. (A) Drops (10  $\mu$ L) of water or 0.2% SDS were placed on the colony surface of each strain grown on oatmeal agar plates for 10 d; (B) Quantitative measurement of *MPG1* and *MHP1* gene expression in the mycelia of  $\Delta$ *Mopas1* by qRT-PCR, normalized to  $\beta$ -tubulin and expressed relative to expression in the mycelia of the wild-type strain.

and hyphae [29]. We next examined surface hydrophobicity by dropping water or SDS on the colony surface of the wild-type strain and  $\Delta$ *Mopas1* (Figure 5(A)). When water drops were placed on wild-type and  $\Delta$ *Mopas1*, the drops were suspended on the aerial surface of both strains (Figure 5(A)). However, when the detergent agent SDS was dropped onto the strains, SDS drops immediately soaked into the aerial structures of  $\Delta$ *Mopas1* but remained on the surface of the wild-type strain. To investigate whether the defect of hydrophobicity in  $\Delta$ *Mopas1* was related to hydrophobins, we measured the mRNA levels of hydrophobin genes *MPG1* and *MHP1* in  $\Delta$ *Mopas1* using qRT-PCR (Figure 5(B)). *MPG1* expression increased in  $\Delta$ *Mopas1* compared with the wild-type, whereas *MHP1* expression decreased (Figure 5(B)). This result suggests that *MoPAS1*-mediated regulation is involved in *MPG1* induction and *MHP1* repression to enable hydrophobicity in fungal tissue.

### 3.5. *MoPAS1* is important for fungal pathogenicity

To determine fungal pathogenicity, rice leaves were inoculated using mycelial agar plugs grown on OMA (Figure 6(A)). Wild-type developed disease lesions on rice leaves, but  $\Delta$ *Mopas1* failed to generate disease lesions (Figure 6(A)). We repeated this experiment with leaves wounded by puncture using a needle. On wounded leaves, mutants formed disease lesions similar to the wild-type (Figure 6(A)). When conidial suspensions of strains were sprayed on rice leaves, wild-type created disease lesions, whereas  $\Delta$ *Mopas1* did not (Figure 6(B)). To investigate whether this loss of pathogenicity was due to appressorium-mediated penetration, we performed penetration assays of rice sheath and onion epidermis (Figure 6(C)). In both

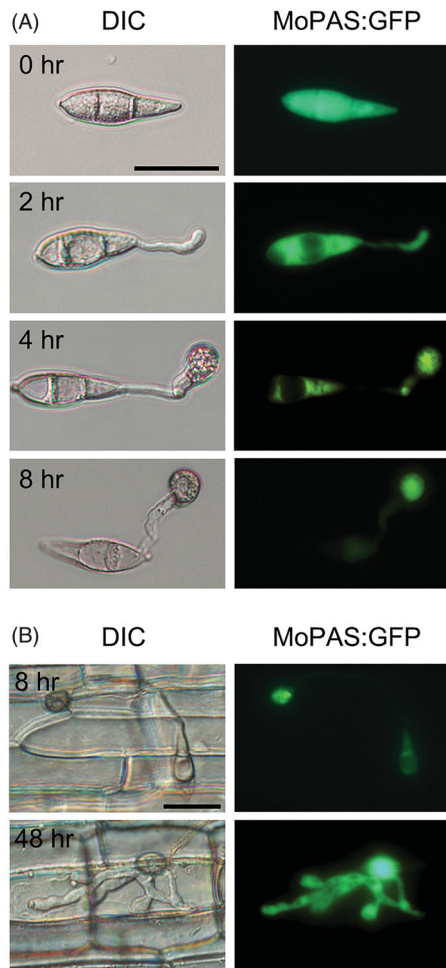


**Figure 6.** Pathogenicity assays. (A) Leaves of the rice cultivar Nakdongbyeo were inoculated with mycelial agar plugs and incubated for 7 d; (B) Conidial suspensions were sprayed onto rice leaves and incubated for 7 d; (C) Conidial suspensions of each strain were inserted into rice sheath and onion epidermis and incubated for 48 h. Bar = 50  $\mu$ m.

rice sheath and onion epidermis, abnormal appressoria were observed when  $\Delta$ *Mopas1* was inoculated, and most of these abnormal appressoria formed could not penetrate the rice sheath or onion epidermis. The invasive hyphal growth was also significantly attenuated compared to the wild-type (Figure 6(C)). In contrast, the wild-type and *Mopas1c* developed normal appressoria and were able to penetrate the plants. These results suggest that *MoPAS1* plays an important role in appressorium-mediated penetration in *M. oryzae*.

### 3.6. Subcellular localization of *MoPAS1* protein

To investigate the localization of *MoPAS1*, we constructed *MoPAS1*:GFP and inserted it into the protoplasts of wild-type. On hydrophobic cover glasses, which induces appressorium formation of *M. oryzae*, the *MoPAS1*:GFP fusion proteins were initially distributed uniformly in the cytoplasm of conidia (Figure 7(A)). As conidia germinates, the *MoPAS1*:GFP signal was observed in the germ tube. After 8 h, *MoPAS1*:GFP fusion proteins in conidia and germ tubes disappeared and were only observed inside appressoria. On rice sheath cells, *MoPAS1*:GFP fusion proteins were intensely localized inside appressoria after 8 h, and uniformly distributed in intracellular invasive hyphae (Figure 7(B)).



**Figure 7.** Localization of MoPAS1:GFP in *Magnaporthe oryzae* strain KJ201 on hydrophobic coverslips or in plants. (A) During appressorium development on hydrophobic coverslips, MoPAS1:GFP was localized in the conidia, germ tubes, and appressoria; (B) *In planta* observations in rice sheath. MoPAS1:GFP were uniformly localized in invasive hypha. Bar = 20 μm.

#### 4. Discussion

We searched for PAS domains in a fungal database (fungiDB.org) and identified a total of 14 genes containing PAS domains in the *M. oryzae* strain KJ201 genome. Among these genes, six were identified to contain HisKA (signal transduction HK) domains (Figure 1). HK is a sensor protein that is typically found in fungi as a hybrid protein including HisKA and REC (signal receiver) domains [30]. It has been reported that PAS-containing HKs are important for infection in plant-pathogenic fungi [18]. Other genes also contained signal transduction domains, including RGS (regulator of G protein signaling) and S/T (serine/threonine kinase) domains. The RGS domain is a conserved domain found in G proteins that regulates G protein-mediated cell signaling [31,32]. For example, in fungi, GPCRs trigger signaling cascades such as the cyclic AMP-dependent protein kinase (cAMP-PKA) or mitogen-activated protein kinase (MAPK) cascades, leading to the

activation of cellular processes such as vegetative growth, morphogenesis, and appressorium formation [33]. These results suggest the importance of PAS domains as diverse regulators of cellular functions in fungi.

In the current study, we investigated the functional roles of *MoPAS1* (MGG\_02665) using a targeted gene-deletion strategy.  $\Delta MoPAS1$  did not form appressoria on hydrophobic surfaces, but this defect was restored when 5 mM cAMP was added to  $\Delta MoPAS1$  (Figure 4). A previous study revealed the importance of cAMP in the infection-related development of *M. oryzae* [34], and found that the addition of cAMP induced appressoria formation by *M. oryzae* on a non-inductive surface. Another study revealed that a protein kinase, *MoYAK1*, is associated with the cAMP pathway [25]. Consistent with the results of the present study,  $\Delta MoYAK1$  did not form appressoria on hydrophobic surfaces, but the addition of cAMP induced appressoria formation. Together, the results of the current study and previous reports suggest that the defect in preinfection-associated developmental stages in *M. oryzae* may be related to the signaling pathway of cAMP-mediated appressorium formation.

$\Delta MoPAS1$  produced significantly fewer lesions on rice leaves, and the conidia of  $\Delta MoPAS1$  formed irregularly shaped appressoria on rice sheath and onion epidermis (Figure 6). A recent study highlighted the importance of HKs in fungal pathogenicity in *M. oryzae* [18]. In this previous study, 10 putative HK-encoding genes were functionally characterized in *M. oryzae* strain 70-15. In these 10 putative genes, the PAS-containing proteins Hik2p, Hik4p, Hik5p, and Hik8p were involved in pathogenicity on rice cultivar CO-39. However, another PAS-containing protein, Hik9p, was not involved in pathogenicity.  $\Delta Mohik9$  (MGG\_02665) caused disease lesions in the rice cultivar CO-39 similar to those of the wild-type 70-15 strain. These results differed from those of the present study, which may be attributable to the different wild-type strains of *M. oryzae* used, or to differences among rice cultivars. We found that MoPAS1:GFP fusion protein was localized in the cytosol of conidia, germ tubes, and appressoria during appressorium development on a hydrophobic coverslip. The fusion protein was expressed during plant infection and intensely concentrated in the penetrating appressorium (Figure 7(A,B)). These results support the conclusion that MoPAS1 plays an important role in disease development in *M. oryzae* KJ201.

Hydrophobins confer surface hydrophobicity to fungal conidia and hyphae by coating external surfaces [29]. The mycelial surface of  $\Delta MoPAS1$  was detergent-wettable, unlike that of the wild-type, and



the relative expression levels of the hydrophobin-containing genes *MPG1* and *MHP1* were also altered compared to those of the wild-type (Figure 5). It has been reported that the deletion of *MHP1* affects conidiation, appressorium development, and pathogenicity [35]. It has also been reported that *MoYAK1*, an *M. oryzae* protein kinase gene, is involved in the regulation of surface hydrophobicity [25]. In  $\Delta$ *Moyak1*, the expression of *MPG1* and *MHP1* was significantly altered.  $\Delta$ *Moyak1* exhibited an easily wettable phenotype and reduced conidiation compared to the wild-type, as well as defective appressorium formation. The defect in appressorium formation was restored by the addition of 5 mM cAMP. These results suggest that *MoPAS1* is involved in the regulation of *MPG1* and *MHP1* and plays an important role in surface hydrophobicity for *M. oryzae*. We suggest that *MoPAS1* is involved in various cellular functions associated with conidiation, surface hydrophobicity, and disease development in *M. oryzae* strain KJ201. Further studies on the other members of the PAS domain family will help to characterize fungal development and pathogenicity in response to diverse environmental signals.

### Disclosure statement

No potential conflict of interest was reported by the authors.

### Funding

This study was supported by Basic Science Research Program through the National Research Foundation of Korea grant [NRF-2017R1D1A1B03029622] funded by the Ministry of Education, Science and Technology, and by a grant [918019-04] from the Strategic Initiative for Microbiomes in Agriculture and Food, Ministry of Agriculture, Food, and Rural Affairs, Republic of Korea.

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