Regulation of Cellular Diacylglycerol through Lipid Phosphate Phosphatases Is Required for Pathogenesis of the Rice Blast Fungus, *Magnaporthe oryzae*

Md. Abu Sadat1, Junhyun Jeon1, Albely Afifa Mir1, Jaeyoung Choi1, Jaehyuk Choi2, Yong-Hwan Lee1,2,3*

1 Department of Agricultural Biotechnology, Plant Genomics and Breeding Institute, and Research Institute for Agriculture and Life Sciences, Seoul National University, Seoul, Korea, 2 Center for Fungal Pathogenesis, Plant Genomics and Breeding Institute, and Research Institute for Agriculture and Life Sciences, Seoul National University, Seoul, Korea, 3 Center for Fungal Genetic Resources, Plant Genomics and Breeding Institute, and Research Institute for Agriculture and Life Sciences, Seoul National University, Seoul, Korea

Abstract

Considering implication of diacylglycerol in both metabolism and signaling pathways, maintaining proper levels of diacylglycerol (DAG) is critical to cellular homeostasis and development. Except the PIP2-PLC mediated pathway, metabolic pathways leading to generation of DAG converge on dephosphorylation of phosphatidic acid catalyzed by lipid phosphate phosphatases. Here we report the role of such enzymes in a model plant pathogenic fungus, *Magnaporthe oryzae*. We identified five genes encoding putative lipid phosphate phosphatases (*MoLPP1* to *MoLPP5*). Targeted disruption of four genes (except *MoLPP4*) showed that *MoLPP3* and *MoLPP5* are required for normal progression of infection-specific development and proliferation within host plants, whereas *MoLPP1* and *MoLPP2* are indispensable for fungal pathogenicity. Reintroduction of *MoLPP3* and *MoLPP5* into individual deletion mutants restored all the defects. Furthermore, exogenous addition of saturated DAG not only restored defect in appressorium formation but also complemented reduced virulence in both mutants. Taken together, our data indicate differential roles of lipid phosphate phosphatase genes and requirement of proper regulation of cellular DAGs for fungal development and pathogenesis.

Introduction

Diacylglycerol (DAG) plays crucial roles in cells as a second messenger in lipid-mediated signaling pathway and as an intermediate in lipid metabolism [1,2]. DAGs is not a single molecular species but a pool of molecules varying with acyl chain length and saturation level [3,4]. Mammalian cells produce more than 50 different types of DAGs including polyunsaturated, diunsaturated, monounsaturated or saturated forms [5]. Different DAGs interact with a diverse array of proteins with C1 domain(s) having different specificities and affinities for DAG, leading to remarkable complexity in DAG-dependent cellular processes [6].

Yeast and mammals have two *de novo* pathways for production of DAG [7]. In one pathway, DAG is synthesized from glycerol-3-phosphate and in another pathway, DAG is generated from dihydroxyacetone phosphate. These two precursors produce lysophosphatidic acid (LPA) and phosphatidic acid (PA) through two acylation steps and finally PA is transformed to DAG by the action of lipid phosphate phosphatase (LPP) [8] (Figure S1). In addition to *de novo* pathways, DAGs can be produced in a manner that is highly dependent on extracellular stimulation. Polyunsaturated DAG is generated from phosphatidyl-inositol-4,5-bisphosphate (PIP2) by the activity of phospholipase C (PLC) through a single step reaction [9]. Alternatively, monounsaturated/saturated DAGs can be generated in a two-step reaction. In the first step, monounsaturated/saturated phosphatic acid (PA) is produced from phospholipids through the activity of PLD and in the second step, dephosphorylation of PA by the members of LPP family enzyme results in monounsaturated/saturated DAG [10]. Yeast has two different types of the enzymes, lipid phosphate phosphatase (LPP) and diacylglycerol pyrophosphate phosphatase (DPP) [10] to dephosphorylate PA, whereas mammals lack DPP. Both the LPP and DPP are the members of LPP family.

All the pathways except the one involving PLC converge on dephosphorylation reaction of PA, indicating the importance of LPP in lipid metabolism and DAG-mediated signaling pathways. Due to its status as a gateway to DAG production, LPPs have been
well studied and documented in diverse organisms ranging from yeast to plant and insects. In Arabidopsis thaliana, AtLPP1 and AtLPP2 are involved in stress response and regulation of stomatal movement through ABA signaling, respectively [11,12]. LPP is shown to play important roles in germ cell migration and tracheal development in insect [13,14]. In yeast, deletion of individual or both lipid phosphate phosphatase (SlLPP1) and diacylglycerol pyrophosphate phosphatase (SdLPP1) did not show any visible phenotypes compared to wild type strain, but both genes together controlled cellular reservoir of the phosphatidic acid (PA), lysophosphatidic acid (LPA) and diacylglycerol pyrophosphate (DGPP) [10,15]. Despite the importance of LPP-mediated regulation of cellular DAGs in different organisms, its implication in fungal pathogenesis remains unexplored. Here we set out to investigate the role of LPP encoding enzymes in development and pathogenesis of a model plant pathogenic fungus, Magnaporthe oryzae.

M. oryzae is a filamentous fungus that causes the rice blast disease. The rice blast disease is one of the most devastating fungal diseases of rice throughout the world [16]. This disease causes 11 – 30% yield losses of the world rice production and is responsible for recurring epidemics throughout South East Asia and South America [17]. Infection by this fungus begins when an asexual spore called conidium germinates following tight adherence to the surface of host plants. Upon recognition of environmental cues such as surface hydrophobicity, the tip of germ tube develops into a dome-shaped, specialized infection structure called an appressorium [18]. Using turgor pressure generated in appressorium, the fungus mechanically rupture the cuticular layer of the plant and gain access to host tissues [19]. Once inside the host cells, the fungus develops ramifying bulbous invasive hyphae that actively grow to result in visible disease lesion, from which massive number of conidia are produced as secondary inoculum [20–22].

Due to genetic tractability of rice blast pathogen and rice [23–25], signaling transduction pathways involving cAMP, calcium, and MAP kinase have been well documented for infection-related development in this fungus [26–31]. Moreover, other studies suggested that DAG plays significant role in pathogenicity leading to appressorium formation in M. oryzae [32,33]. Although a number of studies suggested that lipids and their intermediates are implicated in pathogenesis of fungi including M. oryzae [34–40], genetic analysis on regulation of cellular DAG, a key element of lipid metabolism and signaling has not been carried out.

As the first step to elucidate the biological functions of LPP genes in M. oryzae, we have identified genes encoding LPP family and functionally characterized those genes through gene deletion approach. Analyses of the deletion mutants showed that individual deletion of MoLPP3 and MoLPP5 caused defect in appressorium formation and pathogenicity. Exogenous addition of saturated DAG restored both appressorium formation and virulence defect in both mutants, indicating that maintaining DAG homeostasis is required for fungal pathogenesis. Our work shed light on the critical roles of lipid metabolism during fungal pathogenesis.

Results

Identification of LPP genes in M. oryzae

To identify the members of PAP2 domain containing LPP encoding genes involved in DAG metabolism in M. oryzae, we searched the genome of M. oryzae for genes encoding phosphatidic acid phosphatase type 2/haloperoxidase (PAP2) domain (IPR000326), which is a signature of LPP enzymes [http://cbgp. riceblast.snu.ac.kr] [41]. Our search found a total of eight genes encoding PAP2 domain. BLAST search using amino acid sequences encoded by these eight genes as queries showed that five of them were homologous to yeast LPP1 (YDR503C) and DPP1 (YDR284C) (MoLPP1; MGG_09330.6, MoLPP2; MGG_09380.6, MoLPP3; MGG_09994.6, MoLPP4; MGG_05630.6, MoLPP5; MGG_12462.6), and the rest of them to vanadium chloroperoxidase (MoVAN; MGG_02210.6), long chain base protein 3 (MoLCBP3; MGG_17305.6) and dolichylphosphate phosphatase (MoDpp; MGG_09194.6), respectively (Figure S2). Multiple sequence alignment and hydropathy plot revealed that the MoLPP1, unlike the rest of PAP2 domain-containing proteins, have conserved three sequence motifs (KXXXXXXR, PSPH4 and SRXXXHHXXXXD) (Figure 1A) [42] and six transmembrane domains (Figure 1B and Figure S3), both of which are common features of LPP and DPP enzymes [43]. Based on this data, we only focused on MoLPP1 to MoLPP5 in the following analyses. De novo synthesis of DAG is known to occur mainly in the endoplasmic reticulum (ER). In accordance with this, PSORT predicted that all MoLPPs are localized to ER [44].

Expression analysis of MoLPP genes

As an attempt to infer the implication of MoLPP genes in M. oryzae, qRT-PCR was done with cDNAs sampled from different developmental stages. Expression profiling revealed that the transcript abundance of genes correlates with each other except with MoLPP4 (Figure 2). Correlation coefficient of MoLPP4 with expression of other MoLPP genes ranged from 0.29 to 0.73, while correlation coefficient for all pairs of other genes was higher than 0.75. This suggests that MoLPP genes except MoLPP4 are likely to share overlapping regulatory mechanisms for transcription. In comparison with germinating conidia, most genes were down-regulated in conidia and appressoria stages. But interestingly, MoLPP4 was the only up-regulated gene found in appressorium. However, they tended to be up-regulated during host infection [45]. Based on this, the expression of MoLPP2 and MoLPP5 remained relatively high, compared to the others (Figure 2). In consistent with our observation, the expression of MoLPP2 and MoLPP5 genes were detected in in planta EST library [45]. We also checked the expression levels of MoLPP genes at 40–45 hpi (hours post inoculation) through RNA-Seq data (unpublished data). We found up-regulated expression of MoLPP2 and MoLPP5 compared to mycelial stage, whereas the rest three genes were down-regulated at the same stage. Our data suggest the possible involvement of MoLPP genes in post-penetration phase of host infection by the rice blast fungus.

Targeted gene disruption of MoLPPs

To investigate the roles of MoLPP genes in M. oryzae, we generated deletion mutants of individual genes. Knockout constructs were prepared via double joint PCR [46] and directly used for transformation of wild-type protoplasts. Correct gene replacement event in the resulting transformants was confirmed by PCR-based screening and subsequent Southern hybridization analysis (Figure S3). The deletion mutants were obtained for all but one gene. Despite our repeated efforts, we were not able to generate a deletion mutant for MoLPP4. Consequently, we analyzed four genes, of which deletion mutants are available.

Vegetative growth, germination, and appressorium formation of the mutants

During phenotype analysis, we found that none of the MoLPPs were required for vegetative growth and asexual reproduction (Table 1). When the suspensions of mutant spores were incubated on inductive surface, spores of all mutants were capable of germinating and developing appressoria. However, we noticed that both germination and appressorium formation were delayed.
in ΔMoLpP3 and only - appressorium formation was delayed in ΔMoLpP5 (Figure 3B). Despite - delayed appressorium formation by the two mutants, morphology of appressoria was indistinguishable from that of wild-type. Such delay in germination and appressorium formation in the mutants was complemented by introducing the sequences of *MoLPP3* and *MoLPP5* genes back into the genome of each deletion mutant (Figure 3B).

**Requirement of *MoLPP3* and *MoLPP5* for full virulence of the fungus**

Next, we asked if *MoLPP* genes are involved in fungal pathogenesis on host plants as suggested by the expression analysis. To test this, conidial suspensions of the mutants were spray-inoculated onto rice plants of a susceptible cultivar, Nakdongbyeo. Pathogenicity test showed that ΔMoLpP1 and ΔMoLpP2 were comparable to the wild type in their ability to cause disease on rice plants, whereas ΔMoLpP3 and ΔMoLpP5 were
not able to produce as large and many number of disease lesions on rice leaves as the wild-type (Figure 4A). Measurable and significant reduction in number and size of disease lesions were observed for the two mutants, resulting in 62 to 65% reduction in diseased leaf area (DLA), compared to the DLA calculated for leaves inoculated with wild-type strain (Figure 4B). The complementation strains of ΔMolpp3 and ΔMolpp5 were as virulent as the wild-type, indicating that Molpp3 and Molpp5 are required for full virulence of the fungus.

Defect in virulence observed for ΔMolpp3 and ΔMolpp5 could not be accounted for by minor delay in appressorium formation. Therefore, we monitored early infection process using sheath inoculation method [47]. The mutant appressoria were able to penetrate plant cells as efficiently as the wild-type appressoria. However, invasive hyphae of both ΔMolpp3 and ΔMolpp5 were largely restricted to the primary infected cells at 48 hour post inoculation (hpi) in contrast to invasive hyphae of wild type actively growing to fill in the first host cell and moving to neighboring cells. (Figure 4C and 4D). This result indicates that Molpp3 and Molpp5 are necessary for the fungus to grow inside host plants in early infection stage.

To date, one of the major causes of defective invasive growth is sensitivity to or inability of the fungus to scavenge/suppresses reactive oxygen species (ROS) that are produced by host plants as a defense response during host-pathogen interaction [48,49]. In order to test this possibility, we examined sensitivity of the fungus to oxidative stress in the form of H2O2 or methyl viologen (MV). To test such imbalance in the mutants, we exogenously added saturated DAG (1,2-dicotnly-sn-glycerol: DOG, Sigma-Aldrich) in the conidial suspension and checked appressorium formation at different time points. We also checked appressorium formation after adding CaCl2.2H2O alone or combination of DAG and CaCl2.2H2O to examine whether calcium signaling pathway is implicated. We found that addition of saturated DAG was able to restore appressorium formation without delay in the mutants, while CaCl2.2H2O had little effect in isolation or combination with DAG (Figure 3A). More importantly, when conidial suspensions of the mutants supplemented with DAG were used for spray-inoculation onto rice plants, virulence of both ΔMolpp3 and ΔMolpp5 were recovered to the wild-type level (Figure 3B). These data suggest that Molpp3 and Molpp5 are involved in regulation of cellular DAG and such regulation is important for fungal pathogenesis.

Transcriptional expression pattern of genes involved in cellular DAG production

Given the different phenotypic consequences of deletion of Molpp genes, we investigated the relationship among the genes by checking the transcript abundance of genes in each deletion mutant background. Our transcript analysis showed that deletion of Molpp1 or Molpp2 increased the expression of rest of the genes, whereas deletion of Molpp5 or Molpp5 decreased the expression of other genes (Figure 6). It appeared that deletion of Molpp1 or Molpp2 can be complemented by elevating the expression of Molpp3 and/or Molpp5 but the reverse could not be achieved. These data suggest that transcriptional regulations of LPP genes are intertwined in a network where expressions of other genes are dependent on expression of Molpp3 or Molpp5.

Although PI-PLC-mediated pathway is known to produce different type of DAGs in mammals, we could not exclude the possibility that this pathway contributes to maintaining homeostasis by responding to perturbation in cellular reservoir of DAG. However, when we checked the expression of three PLC encoded genes (MolPLC1, MolPLC2 and MolPLC3) in ΔMolpp3 and ΔMolpp5 background, none of them showed significant difference relative to the wild type strain, suggesting that Molpp-dependent and PI-PLC-mediated pathways are under independent regulation (Figure S5). Therefore, we rule out the possibility that depletion of cellular DAG level resulted from deletion of LPP genes can be complemented by up-regulation of the alternative pathway.

Discussion

In many pathogenic fungi, it has been shown that DAG acts as a second messenger involved in regulation of developmental processes as a part of PLC-mediated signaling pathways [29,50–52]. However, roles of DAG are not limited to a second messenger but as diverse as a basic component of membranes, a precursor in lipid metabolism and a central element in lipid-mediated signaling pathway [53]. Such a broad implication of DAG in cellular processes implies that cellular DAG level should be tightly regulated. A critical point at which DAG production and clearance are regulated is the dephosphorylation of PA by LPP.

Here we identified LPP encoding genes using three criteria: presence of PAP2 domain, conservation of three sequence motifs, and presence of six trans-membrane domains in gene-coded proteins. Among the eight genes encoding PAP2 domain, the five that meet the rest of criteria, were selected as candidate LPP genes in M. oryzae and named Molpp1 to Molpp5. The fact that three distinct motifs and six trans-membrane domains are the hallmark properties of LPP enzymes, suggesting that Molpp1 to Molpp5 are likely to be bona fide LPPs. It is not clear whether or not

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**Figure 2. Transcript abundance of Molpp genes in development stages in Magnaporthe oryzae.** My, mycelia; Con, germinating conidia; App, appressoria; IF 72, early infection stage at 72 hour post inoculation (hpi); IF 148, late infection stage at 148 hour post inoculation (hpi).

Figure 2. Transcript abundance of Molpp genes in development stages in Magnaporthe oryzae. My, mycelia; Con, germinating conidia; App, appressoria; IF 72, early infection stage at 72 hour post inoculation (hpi); IF 148, late infection stage at 148 hour post inoculation (hpi). doi:10.1371/journal.pone.0100726.g002
MoVAN, MoLCBP3, and MoDoPP have phosphatase activity, considering poor conservation of three motifs that are important for catalytic activity. Among the five putative LPPs, MoLPP4 and MoLPP5 were more closely related to DPP1 than LPP1 in S. cerevisiae, suggesting that their substrates may include DGPP (Figure S2). Of note, MoLPP4 seemed to be particularly highly divergent from the yeast LPP1. In conjunction with that, the expression profile of MoLPP4 deviating the most from the other genes suggests sub-functionalization of MoLPP4. It is currently difficult to further address the implication of such divergence due to our failure to generate a knockout mutant strain for the gene.

One possibility of the failure in generation of mutant lacking MoLPP4 is lethality of gene deletion. However, based on the studies of yeast LPP1 and DPP1, together with our own work, it is not likely that deletion of individual LPP genes including MoLPP4 is lethal. In M. oryzae, as with the other filamentous fungi, it is known that efficiencies of gene deletion by homologous recombination are generally low and dependent on the locus [54,55]. Therefore, we presume that the genomic environment of the locus harboring MoLPP4 is not favorable for homologous recombination to occur.

Among the remaining four LPP genes, deletion of either MoLPP3 or MoLPP5 resulted in the fungus that was significantly impaired in pathogenesis, in contrast to deletion of MoLPP1 or MoLPP2 having no effects on fungal developments and pathogenicity. This discrepancy in effect of deletion between two sets of...
genes could be explained by our expression analysis that revealed complex regulatory relationship among LPP genes. Since LPPs are not able to directly regulate expression of genes, the nature of such regulatory relationship should be dependent on lipid signaling. Removal of terminal phosphomonoester group from bioactive lipid molecules such as phosphatidic acid through LPP activities is known to result in functional inactivation of these lipids, which otherwise have multitude of effects on cells through interaction with their targets [56]. We conjecture that each LPP might have different but overlapping substrate specificity and that regulation of bioactive lipids by different LPPs would underlie the observed transcriptional network among LPP genes.

Notably, effect of deletion of MoLPP3 or MoLPP5 was specific to plant infection. This may be possibly due to functional divergence among MoLPP genes after duplication events leading to paralogs in the genome. However, it is still possible that MoLPPs function in combinations, making it difficult for us to evaluate their functions independently of other genes via deletion of individual genes. For example, deletion of MoLPP3 together with other MoLPP(s) may reveal roles of MoLPP3 in vegetative growth. Unfortunately, however, introducing more than three targeted mutations into the fungal genome is technically challenging and our efforts to make double deletion mutant for MoLPP3 and MoLPP5 were unsuccessful.

We showed that virulence defect of ΔMolpp3 and ΔMolpp5 are attributed to the reduced ability of the mutant to grow inside host cells. However, both mutants were comparable to the wild-type in sensitivity to reactive oxygen species and a cell wall perturbing agent, leaving an open question of why invasive growth is significantly impaired in the mutants. In a human pathogen, Cryptococcus neoformans, DAG produced by inositol phosphorylceramide (IPC) synthase is known to regulate production of melanin and antiphagocytic protein, two important virulence factors, in at least two ways [52]. First, DAG can bind to C1 domain of Pkc1 leading to increase in Pkc1 activity, which in turn mediates melanin production. The second way is binding of DAG to the transcription factor Atf2 that promote production of antiphagocytic protein 1. Although the genome of M. oryzae does not encode an orthologs of IPC synthase, it may be possible that LPP-mediated production of DAG may have an impact on regulation of some virulence factors in M. oryzae through its interaction with kinases or transcription factors as shown in C. neoformans.

Alternative possibility is related to the effects of LPP activities on the membrane lipid bilayer through production of DAG. Under equilibrium conditions, DAG contributes no significant proportion of cell membranes. However, local and transient accumulation of DAG in the membranes may lead to changes in physical properties of membrane itself and membrane-associated proteins, influencing important cellular processes such as membrane trafficking and exocytosis [57,58]. In addition, production of DAG will alter lipid compositions in the membranes as well, since DAG is a byproduct of dephosphorylation reaction of lipids. In many intracellular pathogens of animals, lipid rafts, sterol and sphingolipid-rich membrane microdomains, have been shown to control their virulence [59]. Even in human pathogenic fungi, potential role of such membrane domain in pathogenesis was alluded. For example, Candida albicans was shown to produce many GPI-anchored virulence factors that are induced along with sterol-rich domain (SRD) [60]. Similarly, it was suggested that membrane localization and release of the virulence factors such as superoxide dismutase and phospholipase B are regulated by special membrane domains in C. neoformans [61]. Our expression data shows specific up-regulation of MoLPP3 and MoLPP5 during interaction with the host plants. Considering the impact of DAG production on the membranes, it is tempting to speculate that in planta specific activities of LPPs may enable the fungus to modulate or remodel physical properties and lipid composition of membranes for successful colonization of rice plants.

In this study, we investigated how regulation of cellular DAG levels is implicated in development and pathogenesis of the plant pathogenic fungus through deletion of genes that encode proteins involved in a key step of DAG biosynthesis. Our results demonstrated that proper regulation of DAG is pivotal for fungal pathogenesis, independently of the PIP2-PLC pathway. In particular, we showed that LPP-mediated DAG production has profound impact on invasive growth, during which the most intimate interaction with the host plant occurs. Currently other cellular targets of DAG than PKC1 are not known in M. oryzae. Furthermore, there are growing bodies of evidences that suggest the importance of membrane functions in fungal pathogenesis. In light of this, we believe that our work will not only illuminate the importance of DAG production in virulence of plant pathogenic fungi but also have many ramifications on studies regarding lipid metabolisms and membrane-associated processes in fungal pathogenesis.

Table 1. Vegetative growth and asexual reproduction of ΔMolpp1, ΔMolpp2, ΔMolpp3 and ΔMolpp5 in M. oryzae with complementation.

| Strain | Mycelial growth* | Conidiationb |
|--------|------------------|---------------|
|        | CM (mm)          | MM (mm)       | (×10⁷/ml)    |
| K201   | 75.8±1.3         | 70.3±0.6      | 33.4±5.3     |
| ΔMolpp1| 76.3±0.6         | 70.7±0.8      | 32.2±5.6     |
| ΔMolpp2| 76.8±0.3         | 70.3±1.5      | 33.3±6.3     |
| ΔMolpp3| 76.3±1.5         | 70.2±0.8      | 35.2±5.2     |
| ΔMolpp3| 76.8±0.8         | 71.1±1.0      | 35.3±0.3     |
| ΔMolpp5| 76.2±0.3         | 69.5±0.9      | 31.2±2.9     |
| Molpp3c| 75.7±0.6         | 71.5±0.5      | 32.3±3.5     |

*Vegetative growth was measured at 12 dpi on complete agar medium and minimal agar medium. Data were presented as mean ± sd from three independent experiments.

bConidia was measured as the number of conidia from the culture flooded with 5 ml of sterilized distilled water. Data were presented as mean ± sd from three independent experiments.

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Materials and Methods

Fungal isolates and culture conditions

*M. oryzae* wild type strain KJ201 was obtained from the Centre for Fungal Genetic Resources (CFGR: http://cfgr.snu.ac.kr) for this study. All strains used in this study were grown on V8 agar [V8; 8% V8 juice (v/v), 1.5% agar (w/v), adjusted to pH 6.0 using NaOH] or oatmeal agar [OMA; 5% oatmeal (w/v), 2% agar (w/v)] at 25°C in constant light to promote conidial production [62]. All strains cultured on V8 juice agar media for 7 days or on OMA for 10 to 15 days at 25°C under the continuous light condition to observe the developmental and morphologic phenotypes and pathogenicity ability.

Nucleic acids manipulation and expression analysis

Fungal genomic DNA was isolated for two purposes using two different methods. For Southern DNA hybridization, genomic DNA was isolated from mycelia according to a standard protocol [63]. For PCR-based screening of transformants, genomic DNA

Figure 4. Pathogenicity assay of knockout mutants. (A) Disease development after spraying conidial suspension onto rice leaves. Conidia (1 × 10⁵/ml) was sprayed onto the leaves and incubated for 7 days. (B) Comparison of diseased leaf area (DLA). The DLA was calculated relative to the total leaf area using the Axiovision image analyzer. (C) Invasive hyphal growth through sheath assay. Rice sheath was injected with 2 × 10⁶ conidia/ml and observed under microscope at 48 hour post inoculation (hpi). (D) Quantification of invasive growth. Number of cells that are invaded by the fungus was counted using sheath inoculation at 48 hour post inoculation (hpi).

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was extracted by quick and easy method [64]. Southern DNA hybridization was performed on selected transformants to ensure correct gene replacement events and absence of ectopic integration. Genomic DNAs were digested with PstI, BamHI and blots were probed with 1-kb 5'-flanking/3' flanking sequences (Figure 3A). Southern DNA hybridization was done through a

Figure 5. Restoration of appressorium formation and pathogenicity by addition of DAG. (A) Appressorium formation of wild-type and mutants in the presence of DAG and/or CaCl2. Conidial suspension (2×10^4 conidia/ml) was placed on the hydrophobic side of cover slips and mixed with DAG and/or CaCl2 to final concentrations of 20 μg/ml and 10 mM, respectively. Appressorium formation was observed under a microscope 8 and 12 h after incubation. (B) Spray inoculation (upper panels) and sheath assay (lower panels) with conidial suspensions supplemented with 20 μg/ml of DAG.
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To perform the expression analysis by quantitative real-time PCR (qRT-PCR), cDNA synthesis was performed with 5 μg of total RNA using the oligo dT primer with the ImProm-II™ Reverse Transcription System kit (Promega, Madison, WI, USA) following the manufacturer’s instruction. Primer pairs used are listed in Table S1. To compare the relative abundance of MoLPPs (1–5) transcripts, average threshold cycle (Ct) was normalized to that of β-tubulin gene and germinating conidia for each sample.

**Targeted deletion of genes and complementation experiments**

Targeted deletions of genes were carried out by transforming wild-type protoplast with knockout constructs. Knockout construct of individual genes were obtained by double-joint PCR where ~1 kb flanking sequences of target gene were amplified and fused to hygromycin cassette. The resulting transformants were subjected to PCR-based screening and gene replacement event in the genome of transformants were confirmed by Southern blot analysis using one of flanking sequences as a probe. Complemented strains for two mutants (ΔMolpp3 and ΔMolpp5) were generated by reintroducing 1.2 kb and 0.9 kb fragments carrying the MoLPP3 and MoLPP5 ORF, respectively. Complemented strains were selected through PCR screening with specific primers (MoLPP3_NF, MoLPP3_NR, MoLPP5_NF and MoLPP5_NR) (Table S1) and also used “c” to indicate complementation strain through the whole manuscript.

**Developmental phenotype assay**

Radial mycelial growth rate with three replications was measured on complete agar media and on minimal agar media 12 days after inoculation. For conidial germination and appresorium formation, conidia were harvested from 7-days-old V8 juice agar plates and passed through the two layer of miracloth with sterilized distilled water. Concentration of conidial suspension was adjusted to 2 × 10^6 conidia per milliliter and 40 μl was dropped onto coverslips with three replicates and incubated in moistened box at room temperature. At two and four hours after incubation, germination rate was determined by observing at least a hundred of conidia per replicate under a microscope. The rate of appressorium formation was measured as the percentage of germinated conidia that developed appressoria at 8, 16 and 24 hours each after incubation. These assays were performed with three replicates in three independent experiments. For conidiation, conidia was collected from 7-days-old V8 juice agar plates with 5 ml sterilized distilled water and measured by counting the number of asexual spores within 10 μl conidia suspension onto haemacytometer under a microscope.

**Pathogenicity assay and infectious growth visualization**

For spray inoculation, conidia were collected from 7-days-old V8 juice agar media with 10 ml of filtered conidia suspension and adjusted to 1 × 10^5 conidia per milliliter containing Tween 20 (250 ppm final concentration) and sprayed onto the rice seedlings (Oryza sativa cv. Nakdongbyeo) in three to four leaf stage. Inoculated rice seedlings were placed in a dew chamber for 24 hours under the dark condition at 25°C. After then, they were transferred to the rice growth incubator that is maintained at 25°C, 80% humidity and with photoperiod of 16 hours using fluorescent lights [66]. For microscopic observation of invasive growth on rice tissue, excised rice sheath of Nakdongbyeo were prepared by standard process [47]. Conidia suspension was injected in excised rice sheaths and incubated in moistened box for 24 and 48 hours at room temperature. After incubation, the infected rice sheaths were
trumped to remove chlorophyll enriched plant parts. Remaining epidermal layers of mid vein (three to four cell layers thick) were used for microscopic experiment.

Computational analysis

All sequence information used in this study was taken and analyzed from the online database Comparative Fungal Genomics Platform (CFGP) [41], http://cfgp.snu.ac.kr/ and BLAST program provided at the National Center for Biotechnology Information, Bethesda, USA [http://www.ncbi.nlm.nih.gov/blast/ [67]]. Sequences were aligned by ClustalW algorithm [68]. Hydrophathy plot was generated using TopPred 2 [http://www.sbc.su.se/~erikw/toppred2/] [69].

Tests for sensitivity to reactive oxygen species and a cell wall perturbing agent

Sensitivity to reactive oxygen species was tested by growing fungi on complete media supplemented with H2O2 and methyl viologen. Test for cell wall integrity was carried out by adding a cell wall perturbing agent, Congo red (200 ppm) to complete media that are inoculated with wild-type and mutant strains.

Supporting Information

Figure S1 The enzymatic reactions leading to DAG generation through different pathway. D P, dihydroxyacetone phosphate; D P Acyl, dihydroxyacetone phosphate acyltransferase; 1-Acyl-D P, 1-Acyl-dihydroxyacetone phosphate; 1-Acyl-D P Red, 1-Acyl-dihydroxyacetone phosphate reductase; G-3-p, glycerol-3-phosphate; G-3-p Acyl, glycerol-3-phosphate acyltransferase; Lyso PA; Lyso phosphatidic acid; PC, phosphatidylcholine; PLD, phospholipase D; PA, phosphaticid acid; PIP2, phosphatidylinositol-4,5-bisphosphate; PLC, phospholipase C; DAG, diacylglycerol.

(TIF)

Figure S2 Sequence similarities of PAP2 domain of eight genes with each other and also with the yeast

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