HYR1-Mediated Detoxification of Reactive Oxygen Species Is Required for Full Virulence in the Rice Blast Fungus

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Abstract
During plant-pathogen interactions, the plant may mount several types of defense responses to either block the pathogen completely or ameliorate the amount of disease. Such responses include release of reactive oxygen species (ROS) to attack the pathogen, as well as formation of cell wall appositions (CWAs) to physically block pathogen penetration. A successful pathogen will likely have its own ROS detoxification mechanisms to cope with this inhospitable environment. Here, we report one such candidate mechanism in the rice blast fungus, Magnaporthe oryzae, governed by a gene we refer to as MoHYR1. This gene (MGG_07460) encodes a glutathione peroxidase (GSHPx) domain, and its homologue in yeast was reported to specifically detoxify phospholipid peroxides. To characterize this gene in M. oryzae, we generated a deletion mutant.Δhyr1 which showed growth inhibition with increased amounts of hydrogen peroxide (H2O2). Moreover, we observed that the fungal mutants had a decreased ability to tolerate ROS generated by a susceptible plant, including ROS found associated with CWAs. Ultimately, this resulted in significantly smaller lesion sizes on both barley and rice. In order to determine how this gene interacts with other (ROS) scavenging-related genes in M. oryzae, we compared expression levels of ten genes in mutant versus wild type with and without H2O2. Our results indicated that the HYR1 gene was important for allowing the fungus to tolerate H2O2 in vitro and in planta and that this ability was directly related to fungal virulence.

Introduction
Molecular oxygen, itself relatively nontoxic, is important to most living organisms on this planet. However, its derivatives, reactive oxygen species (ROS), can lead to oxidative destruction of cells [1]. For example, in mammals, ROS can accelerate aging by making holes in DNA, which may result in cancer and other severe diseases [2]. However, animals, plants and fungi have all adapted to use ROS as key signaling molecules [3]. In plants, ROS play a more positive role as a defense mechanism against attacking pathogens, and are often produced as a first line of defense [4]. In the plant-pathogenic fungus, Magnaporthe oryzae, ROS regulation plays important roles in both development and virulence. ROS itself has been shown to accumulate in the developing and mature appressorium, or fungal penetration structure, while the two NADPH oxidases in M. oryzae, NOX1 and NOX2 are required for proper development of appressoria, as well as full virulence [5]. The catalase gene family member, encoded by C1TB, was shown to also be involved in cell wall integrity as well as virulence, as deletion mutants were altered in hyphal, spore and appressorial morphology [6]. Organisms, therefore, must carefully balance the toxic effects of ROS and the need for ROS in cellular signaling.

There are five major types of ROS in plants: superoxide (O2−), hydrogen peroxide (H2O2), hydroxyl radical (OH), nitric oxide (NO), and singlet oxygen (ΔO2). In plant cells, organelles with an intense rate of electron flow or high oxidizing metabolic activity are major sources of ROS generation [7]. These organelles include mitochondria, chloroplasts and peroxisomes. ROS are also generated via enzymatic sources, such as membrane-associated NADPH oxidases, cell wall peroxidases and oxalate oxidases [8].

ROS play a crucial role during plant defense responses. Oxidative bursts have been detected when plant cells are inoculated with biotrophic pathogens [9], semi-biotrophic pathogens [10], necrotrophic pathogens [11], and pathogen elicitors [12]. More recent studies with M. oryzae that causes rice blast disease, demonstrated that rice produces H2O2 shortly after inoculation with a virulent strain of the fungus [13,14]. The toxic effects of ROS can directly kill pathogens, and as a result, pathogens have developed counter measures [5]. The coexistence of hosts and pathogens side-by-side determines that the increase of resistance in a host will be balanced by the change of virulence in a pathogen, and vice versa. A metabolite fingerprint study of three rice cultivars infected by M. oryzae provided evidence for suppression of plant-associated ROS generation during compat-
Author Summary

Reactive oxygen species (ROS) are antimicrobial compounds and also serve as stimulators and products of plant defense reactions. ROS appear to be active in the critical zone where pathogens and plants come in contact. Therefore, understanding the source, the role, and the destination of ROS in each interacting partner will be crucial for understanding the pathogen-host molecular battle. In this study, we focused on one potential fungal mechanism for ameliorating effects of plant-produced ROS during the early stages of infection. Characterizing the MoHYR1 gene from the rice blast fungus Magnaporthe oryzae, suggested that MoHYR1 was involved in overcoming plant defense-generated ROS. The deletion of this gene caused a virulence defect in M. oryzae, which we believe was associated with the mutant’s inability to detoxify plant-generated ROS. Together, our data suggest that HYR1 is a virulence factor in the rice blast pathogen, and its role in virulence was directly related to sensing and managing plant-generated ROS during early infection events. HYR1 is part of a ROS scavenging and sensing pathway that is well characterized in yeast, and our study is the first to examine this important gene in filamentous fungi.

Results

Identification and characterization of a Glutathione peroxidase domain-containing gene in the genome of M. oryzae

As one of the key members during the oxidative stress response, the yeast Saccharomyces cerevisiae Hyr1/YIR037W (formerly termed Gpx3) was reported to be a glutathione-dependent phospholipid peroxidase (PHGpx) that specifically detoxifies phospholipid peroxides [19]. In order to identify the corresponding gene in M. oryzae, we performed a BlastP analysis against the fully sequenced genomic database of M. oryzae housed at the Broad Institute. Using an E-value of 1e-3 returned a single hit located on Supercontig 20, with an accession number of MGG_074606.1. It is 1315 bp long including two introns, with an open reading frame of 783 bp, which encodes a 172-aminio acid protein. A sequence analysis was performed using Prosite on the ExPASy Proteomics Server (http://ca.expasy.org/prosite/). Hits revealed a glutathione peroxidase active site at amino acid positions 27–42, and a glutathione peroxidase signature at amino acid positions 66–73 (Figure 1A).

When a BlastP search was performed against GenBank at NCBI, numerous hits were returned with high similarity scores, from many organisms including fungi and bacteria. An alignment indicates that the putative GSHPx domains of Hyr1 are highly conserved across different organisms (Figure 1B). The MoHyr1 protein shares the highest amino acid conservation with the model, non-pathogenic fungus, Neurospora crassa (93% similarity and 73% identity), but shares between 81 and 90% similarity with eight other plant pathogenic filamentous fungi examined (Table S1 and Figure 1C).

Secondary structure of the HYR1 protein was determined by PSIPRED [27], and consists of eight β-sheets (or strands) and four α-helices (Figure 2). As described in Zhang et al [18], the ScHyrlp showed a typical ‘thioredoxin fold’, also consisting of four β-sheets surrounded by three α-helices [28]. We compared the crystal structure of ScHyrlp with the predicted tertiary structure of MoHyr1 protein, generated with PyMOL (http://www.pymol.org/). The MoHyr1 predicted structure appears similar to a canonical thioredoxin fold, showing four β-sheets, with β1 and β2 running parallel and β3 and β4 running anti-parallel, surrounded by three α-helices (Figure 2). We located three positionally conserved cysteines in our HYR1 protein model compared to yeast, and these are marked in Figures 1B and 2. Two important cysteines, Cys39 and Cys80, likely correspond with two active sites found in the yeast Hyr1p, Cys36 and Cys82. Together, our in silico data suggest that we have identified the structural homolog of the ScHyr1 from yeast, and that this gene is highly conserved across filamentous fungi.

In order to functionally characterize the MoHYR1 gene, we obtained the ATCC S. cerevisiae Abhyr1 mutant and its wild type parent for complementation tests. Our hypothesis was that based on its sequence and predicted tertiary structure, the MoHYR1 gene would rescue the yeast mutant when grown on non-permissive concentrations of hydrogen peroxide. As shown in Figure 3, the yeast mutant and the wild type strain both grow well on 0 and 2 mM H2O2. However, growth of yeast Abhyr1 was significantly hindered in 4 mM H2O2. The wild type MoHYR1 gene was transformed into the yeast mutant, which restored partial growth on this higher concentration. To further support our hypothesis, we constructed mutations in the two conserved cysteine residues at positions 39 and 88. Neither of the mutations rescued the yeast phenotype on hydrogen peroxide (Figure 3).
Targeted deletion of MoHYR1

To explore the biological role of the MoHYR1 protein in the development and pathogenicity processes of *M. oryzae*, the deletion mutant *Dhyr1* was generated through homologous recombination of the MoHYR1 open reading frame with a gene conferring hygromycin resistance (hygromycin phosphotransferase; HPH) (Figure S1A). A gene deletion fragment was generated by nested PCR amplification of the 5’ flanking region of MoHYR1, the HPH gene, and 3’ flanking region of MoHYR1, using adapters to link the three pieces together. This gene deletion fragment, which contained flanking regions homologous to the MoHYR1 gene, was introduced into protoplasts of *M. oryzae* via PEG-mediated fungal transformation. After PCR screening of successful knockouts and ectopics using primer pairs outside the flanking regions and inside the HPH gene, two *Dhyr1* knockout mutants (B25, B33) and two ectopic mutants (B40, B60) were identified (Figure S1B) and confirmed with Southerns (Figure S1C). Real-time qRT-PCR was also employed to confirm full deletion of the MoHYR1 gene and no transcripts were detected. Deletion mutant *Dhyr1* (B33) was complemented with a full-length copy of the MoHYR1 gene linked to the cerulean fluorescent protein (Figure S1D, see Materials and Methods).

MoHYR1 is required for vegetative hyphal growth in a ROS-rich environment

HYR1p in yeast was reported to not only be a sensor of ROS, but to have scavenging properties as well [19]. To investigate the role of MoHYR1 in scavenging H2O2 during vegetative hyphal growth, we inoculated the same amount of initial mycelia into complete media (CM) containing 0, 5 and 10 mM H2O2. No significant differences were detected among wild type, the *Dhyr1* knockout mutants and the ectopics when growing in 0 mM H2O2. However, the mycelial growth of the *Dhyr1* knockout mutants was severely and significantly affected at 10 mM H2O2 (Figure S2A and B). By contrast, the wild type and ectopics did not display much difference in mycelial growth at any concentration. The complemented mutant line grew slightly better than wild type in all concentrations of H2O2, and upon Southern analysis, we found that four copies had inserted into the genome (Figure S1E). Together, these data indicated that MoHYR1 was responsible for the H2O2 growth tolerance phenotype.

The MoHYR1 gene contributes to virulence in *M. oryzae*

To determine the role of MoHYR1 in virulence, we drop-inoculated detached leaves of three week-old blast-susceptible
MoHYR1 was required for infection-related activities in planta. A fundamental question we wanted to assess was whether MoHYR1 is involved in coping with ROS generated during infection but not for internal ROS levels. We examined whether ROS was present during any of these processes, particularly during initial infection events, or loses the ability to effectively cope with it. We observed that the ROS "haloes" around the appressoria usually localized directly underneath the appressoria (Figure 6). Previous studies have suggested that reactive oxygen species (ROS) are involved in disease development in rice blast (M. oryzae), and that MoHYR1 might play a role in modulating ROS levels. To further investigate this, we analyzed ROS production in planta using a near-infrared fluorophore 9,7'-dichlorodihydrofluorescein diacetate (H2DCFDA) and its reduced form (H2DCF). The results showed that MoHYR1 mutant plants had significantly reduced ROS levels compared to wild type, indicating that MoHYR1 plays a crucial role in modulating ROS levels in planta.

Figure 3. MoHYR1 complements the S. cerevisiae Δhyr1 mutant. The yeast strains BY4741 (wild type) and BY4741 YIR037W (Δhyr1) were obtained from the ATCC. The mutant was complemented with the wild type copy of itself, the MoHYR1 gene, and the MoHYR1 containing mutations at each of the two cysteine residues (cys39Ala and cys88Ala). All strains were spotted onto YPD plates containing 0 mM, 2 mM and 4 mM hydrogen peroxide. Neither the YIR037W strain, nor the two cysteine residue mutants grow at the non-permissive concentration, however the yeast mutant is partially rescued by the MoHYR1 copy. This experiment was repeated ten times with similar results.

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The M. oryzae's disease cycle is initiated when the conidium contacts a hydrophobic surface, inducing it to germinate. The germinated conidium forms a germ tube and appressorium that penetrates the plant surface via turgor pressure and forms a thin penetration peg into the first plant cell [30]. Thus, we first examined whether ROS was present during any of these processes, and if so whether MoHYR1 was involved in coping with it. We inoculated susceptible rice and barley cultivars with the Δhyr1 mutants, ectopics and wild type. ROS was detected using the indicator 2',7'-dichlorodihydrofluorescein diacetate (H2DCFDA) [31]. Conidia of wild type, ectopics and the Δhyr1 mutant all elicited some degree of ROS when inoculated onto barley leaves (Figure 5A–C), whereas ROS was undetectable under the same imaging conditions when non-inoculated leaves were stained (data not shown). The Δhyr1 mutants showed the strongest ROS signal 24 hours post inoculation (hpi) compared to the others. The signal continued in this manner through 48 hours (data not shown). These experiments were repeated six times and the results were consistent across the two independent Δhyr1 mutant lines. ROS signals were quantified via counting the number of 'ROS haloes' found around appressoria and expressing this as a percentage of appressoria counted per sample; a significant difference in signals was observed between the mutants, wild type, and ectopics (Figure 5D). These results indicate that in the absence of the MoHYR1 gene, the fungus can no longer manage the ROS that is generated during initial infection events, or loses the ability to effectively cope with it.

To better understand the reason for reduced virulence in the Δhyr1 mutant, we wished to determine whether internal fungal levels of ROS were altered in the absence of the gene. The deletion mutant and wild type were grown on complete media and stained with nitroblue tetrazolium (NBT) for production of superoxide anions (Figure S3). Results showed little differences between mutant and wild type when examining the entire colony (Figure S3A and F) or aerial hyphae (Figure S3A–D). Fungal internal ROS patterns are different from those generated in planta.

Figure 5C suggested that reactive oxygen species localized mainly around the appressoria. Upon closer inspection, we observed that the ROS "haloes" around the appressoria usually localized directly underneath the appressoria (Figure 6). Previous experiments have shown that the appressoria are the site of infection and that they are sensitive to ROS. MoHYR1 may therefore play a role in modulating ROS levels in the appressoria, which could explain its role in infection-related activities.

MoHYR1 is required for breaking down ROS in planta during infection but not for internal ROS levels. A fundamental question we wanted to assess was whether MoHYR1 was required for infection-related activities in planta.
Figure 4. *Δhyr1* exhibits a virulence defect. *Δhyr1* mutants display a decrease in pathogenicity compared to wild type, on susceptible barley and rice. (A) Conidia of two *Δhyr1* mutants, B25 and B33, were drop-inoculated onto barley cultivar Lacey and show a virulence defect compared to ectopics (B40 and B60), the complemented line (*Δhyr1-C*), or 70-15 (wild type), as manifested by smaller lesions at 7dpi. (B) Quantification of lesion
size reveals a significant difference in virulence between wild type and ectopics, and the mutants. Different letters over the bars indicate a significant difference as determined by a student’s t-test, and a p-value of ≤0.01. (C) Rice plants (cultivar Maratelli) were spray-inoculated with the mutants, ectopics and wild type (as above) and scored for lesion type 7 dpi. (D) Quantification of lesion type (0 = no symptom; 1 = pinhead-sized brown specks; 2 = 1.5 mm brown spots; 3 = 2–3 mm gray spots with brown margins; 4 = many elliptical gray spots longer than 3 mm; 5 = coalesced lesions infecting 50% or more of the leaf area), reveals no difference in lesion types 1-3 however the two mutants do not make any lesion types 4 and 5. Lesions were photographed and measured or scored 7dpi and experiments were repeated twice with similar results. Different letters over the bars indicate a significant difference as determined by a student’s t-test and a p-value of <0.05.

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studies had demonstrated that the rice blast fungus also generates internal ROS during infection-related development, particularly during appressorial maturation and furthermore, that ROS can be secreted from the fungus itself [5]. In order to identify the source of the reactive oxygen species detected in our experiment, we inoculated M. oryzae onto the hydrophobic side of gel-bond, which can mimic the plant surface and induce ROS production in vitro [32]. The result shown in Figure 7 indicated that first, M. oryzae does generate ROS during germ tube and appressorial formation; second, the reactive oxygen species generated by M. oryzae were mostly intracellular and did not appear to be secreted or defined; and finally, that ROS were relatively weak in the fungal structures by 24 hpi. These observations occurred in the wild type, ectopic and mutant lines, indicating little difference in internal ROS levels regardless of the presence of HYR1. Altogether, these results were different from what we observed in planta, which was a strong ROS signal from 24–48 hpi.

Three lines of evidence suggest ROS is most likely plant-generated

In order to identify the source of the ROS detected during susceptible interactions, we used diaminobenzidine (DAB) to study the ROS distribution pattern. barley leaves were inoculated with Ahyr1 mutant then stained with DAB and imaged using confocal reflected light signal to visualize the DAB deposits from a top view of an interaction site (Figure 8A). The leaf samples from this same interaction site was processed further and embedded in epoxy resin to obtain a cross-section using a correlative microscopy approach. The confocal images suggested that the dark region (DAB) was localized immediately adjacent and inside the plant cell wall (Figure 8B) centered around the penetration peg (arrowhead – Figure 8B).

The second piece of evidence resulted from scavenging for ROS with ascorbic acid, an antioxidant that detoxifies hydrogen peroxide [33]. When 0.5 mM ascorbic acid was mixed with Ahyr1 mutant conidia, inoculated onto plants and stained with H$_2$DCFDA, ROS haloes were clearly observed (Figure 9A). However, when barley leaves were pre-treated with ascorbic acid, then inoculated and stained with H$_2$DCFDA, almost no ROS haloes were detected (Figure 9B). This experiment was repeated with another ROS-inhibitor called DPI (diphehyleneiodonium chloride), with similar results (data not shown). Ascorbic acid-treated leaves were also inoculated with mutant conidia and allowed to incubate in the growth chamber for six days, after which time we observed wild type lesions (Figure 9C). This suggested that the ROS haloes observed in this experiment are likely originated from the plant.

Furthermore, we analyzed previously characterized nos1 and nos2 mutants for ROS haloes; in M. oryzae, NOX1 and NOX2 code for NAPDH oxidases, and are largely responsible for producing internal ROS [5]. We hypothesized that if ROS was emanating from the plant, than the loss of the NOX genes should have no effect on haloes. Overall, haloes can still be produced when either of the nos mutants, or its parental strain, Guy11 was inoculated onto barley leaves (Figure S4A–F). While there was a slight significant difference among the number of haloes observed when looking at the individual mutants (nos1 made slightly more than nos2); there was no significant difference between mutants and wild type (20–30 appressoria were counted per strain, and the percentage of those with haloes, reported; Figure S4G).

MoHyr1 has an effect on later, but not immediate, plant-produced ROS

Since our data strongly suggested that Ahyr1 mutants had a lower capacity to eradicate plant-generated ROS during early stages of infection. Our next goal was to determine whether this gene played a role in fungal tolerance to ROS generated immediately following inoculation. In order to carry out this experiment, we inoculated susceptible barley leaves with either the Ahyr1 mutants or the wild type conidia, and imaged them 1 hpi. The ROS dye H$_2$DCFDA was injected directly into the leaves, so the result only showed the redox status inside the leaves, and not inside the fungus, which might have skewed the results. Our data revealed that ROS was detected 1 hpi, which indicated that the plant detected and responded to the pathogen at an early time point (indicated by ROS fluorescence in the mesophyll cells; Figure S5A). A quantitative analysis of the signal intensities by ImageJ (available at http://rsb.info.nih.gov/ij; developed by Wayne Rasband, National Institutes of Health, Bethesda, MD) revealed no significant differences when inoculated with the Ahyr1.

### Table 1. Development characteristics of the Ahyr1 mutant are similar to ectopics and wild type.

| Strain       | Growth rate (cm) | Conidiation$^a$ | % GT$^2$ formation | % AP$^3$ formation | Conidia shape |
|--------------|------------------|-----------------|-------------------|-------------------|---------------|
| 70-15 (WT)  | 5.03±0.32        | 21.33±11.06     | 0.91±0.08         | 0.93±0.06         | normal        |
| Ahyr1 mutant| 5.13±0.06        | 19.1±1.73       | 0.95±0.09         | 0.92±0.02         | normal        |
| Ectopic     | 4.9±0.44         | 20±0            | 0.93±0.08         | 0.97±0.05         | normal        |

$^a$concentration equals 1 × 10$^6$conidia/ml.
$^b$GT = germination tube.
$^c$AP = appressorium.

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mutants or with the wild type conidia (Figure S5B). We thus concluded that the MoHYR1 gene does not play a role in ameliorating an early, or immediate, plant defense response.

To test whether MoHYR1 had any impact on plant-produced ROS that may occur later during infection, we inoculated Dhyr1 mutant conidia or wild type conidia onto barley leaves and stained with DAB at 24 hpi (Figure 10). Results indicated that the Dhyr1 mutant was unable to block ROS produced at 24 hpi, where the ROS was both detected in an entire plant epidermal cell, as well as in plant cells that were not in direct contact with the pathogen (Figure 10).

ROS generated during the infection process are related to cell wall appositions (CWAs)

It has been documented that the presence of reactive oxygen species around CWAs is a biochemical marker for non-penetrated cells during the interaction between barley and barley powdery mildew, Blumeria graminis [34]. To determine whether the ROS observed during a susceptible barley-M. oryzae interaction, we examined leaves that had been inoculated with M. oryzae 24 and 40 hpi with either mutant or wild type conidia. The result showed that classical CWAs were formed within 24 hpi in both strains and no other differences in CWA morphology could be detected (Figure 12).

MoHYR1 regulates other ROS-related genes in M. oryzae

Given the fact that increased ROS accumulation occurs in the absence of MoHYR1, we next tried to determine whether the ROS scavenging system was impaired in the Dhyr1 mutants. We used real-time quantitative real time reverse transcription PCR (real-time qRT-PCR) to compare the expression of general antioxidant and redox control gene orthologs in both M. oryzae wild type and Dhyr1 strains (Figure 13). Primer pairs for the following genes were employed to examine gene expression: YAP1 (MGG_12814.6), GSH1 (g-glutamylcysteine synthetase; MGG_07317.6), GSH2 (glutathione synthetase; MGG_06454.6), GLR1 (glutathione reductase; MGG_12749.6), GTT1 (glutathione transferase 1; MGG_05677.6), SOD1 (Cu/Zn superoxide dismu-
tase; MGG_03350.6), CAT1 (catalase 1; MGG_10061.6), GTO1 (omega class glutathione transferase 1; MGG_05367.6), and cyt c per (cytochrome c peroxidase; MGG_10368.6). The housekeeping gene encoding Ubc (ubiquitin conjugating enzyme; MGG_04081.6) was used as an internal control. We also included the gene MoHYR1 (MGG_07460.6) in this experiment to confirm its deletion in the mutant lines. The expression patterns of these ten genes were placed into two categories. The first category (Figure 13A) is comprised of four genes that show increased expression in the wild type strain after induction with hydrogen peroxide, while expression in the mutant line is low and unchanged. GTI1, GR and GSH1 belong to this category, along with the HYR1 partner protein YAP1; YAP1 also shows slight but significant differences in expression in the Ahyr1 mutant line with and without H₂O₂, and has a higher expression level compared to the wild type strain without ROS. The second category contains genes whose expression does not significantly change, both in response to H₂O₂, as well as in the presence of the MoHYR1 gene. This category includes six genes: cyt c per, CAT1, Cu/Zn SOD, GTI1, GSHII and MoHYR1 (Figure 13B). HYR1 shows no expression at all in the mutant line, which was to be expected.

**Hyr1 cellular localization**

We evaluated the sub-cellular localization pattern of the MoHYR1 protein during infection, conidia of a M. oryzae deletion line (Δhyr1 B33) transformed with cerulean-MoHYR1 N-terminal fusion (the same construct that was used for complementation), was inoculated onto barley leaves and observed during the following time points: 1 hpi, 6 hpi, 12 hpi, 24 hpi and 72hpi. At 1 hpi, MoHYR1 was mainly localized in the conidial vacuoles and with low levels in the cytoplasm. When the germ tube formed, the protein was present throughout the germ tube (Figure 14A). At 6 hpi, the MoHYR1 protein showed increased cytoplasmic localization in the appressorium and conidium and at 12 hours, a concentration of HYR1 in the appressorial cytoplasm (Figures 14B and C). At the later time point, 24 hpi, the protein appeared to be localized in the vacuoles with reduced levels in the cytoplasm (Figure 14D), and a later, invasive stage time point suggests the protein was again cytoplasmically localized (Figure 14E).

**Discussion**

During the interaction between the pathogens and plants, plants mount defense mechanisms to protect themselves from pathogens. The cellular environment within the host can represent a major source of stress towards the invaders [16]. Pathogens, on the other hand, must possess adaptive mechanisms in order to survive. In this study, we hypothesized that the M. oryzae HYR1 protein defines one such mechanism, the glutathione synthesis pathway, involved in coping with the oxidative environment generated by plant defenses.

**MoHYR1 is necessary for ROS detoxification and full virulence**

In M. oryzae, MoHYR1 is the only sequence homolog of the yeast glutathione-dependent peroxidase, HYR1p, formerly termed Gpx3 [35]. In yeast, HYR1p senses H₂O₂ through two highly conserved cysteines that are redox sensitive. Mutations in either of these two cysteines leads to a non-functional HYR1 [18]. Indeed, we found that the wild type MoHYR1, but not the MoHYR1 cysteine

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Figure 6. The ROS observed after inoculation with Δhyr1 conidia as a disk-shaped halo located beneath appressoria. (A) A 3-D projection of confocal images with the ROS stain H₂DCFDA showed a halo (green) of ROS around and beneath the appressoria (blue; AP), which emanated from two nearby conidia. (B) A side-view of panel A showed that the halo was a thin layer of ROS located beneath the appressoria. The ROS halo sits directly between the AP and the plant surface. Scale bar = 10 μm.
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mutants, was able to partially rescue the yeast HYR1p mutant on non-permissive levels of H2O2. This result is similar to *Ayh1* yeast mutants complemented with homologs from two pathogenic filamentous fungi, *Cochliobolus heterostrophus* and *Ustilago maydis*, as both homologs partially complemented the yeast mutation [20,23]. These data suggested that *MoHYR1* may function similarly during redox sensing and the subsequent signaling that leads to ROS detoxification. This model was further supported by the presence of ROS haloes located underneath appressoria during infection with a much greater frequency in the *Ayh1* mutant compared to the wild type strain.

The increase in ROS haloes in *Ayh1* mutants correlated with significantly smaller lesions sizes when inoculated on susceptible rice and barley plants, suggesting that ROS scavenging regulated by *MoHYR1* was required for full virulence. This was supported by a rescuing of the *Ayh1* mutant phenotype to wild type lesions by scavenging plant-derived ROS with ascorbic acid or disrupting plant-derived ROS generation with DPI. These results were similar to a gene recently reported on in the rice blast fungus called DES1 for Defense Suppressor 1 [14]. *DES1* was also involved in virulence and triggers a stronger plant response upon infection, manifested by both an increase of the oxidative burst, as well as expression of two plant defense genes. Intriguingly, *DES1* has no known functional domains and from sequence analysis, its function cannot be predicted, although it is well-conserved throughout fungi. It is also noteworthy that expression of *MoHYR1* was tested in the *Aydes1* mutant, and found to be slightly down-regulated. This could suggest that *HYR1* and *DES1* represent two semi-redundant, semi-dependent mechanisms evolved to cope with the plant defense response. Equally interesting is a gene recently identified in the plant and human fungal pathogens, *Alternaria brassicicola* and *Aspergillus fumigatus*, respectively, called *tmpL* [16]. This membrane-localized gene contains a FAD/NADP-binding domain and had

Figure 7. *Ayh1* (B25) conidia on gel-bond were similar to wild type in terms of ROS production. Staining was performed 24 hpi; Calcofluor White was used to stain the cell walls (blue) and H2DCFDA was used to stain the ROS (green). Conidia of (A) *Ayh1* (B25), (B) wild type (70-15) and (C) ectopic (B40). A transmitted light image was taken as well, and overlaid with the fluorescent image. The inset in panel A showed the fluorescence image of the conidium (1) and appressorium (2). Images were taken using confocal microscopy. Scale bar = 10 μm. doi:10.1371/journal.ppat.1001335.g007

Figure 8. *Ayh1* appressorial-localized ROS appeared to be plant-generated. (A) Reflection confocal imaging with the ROS stain DAB shows a wide ROS signal (arrow) around and beneath the appressorial attachment site (AP). In the middle of the appressorium attachment site was the putative penetration peg site (arrowhead). (B) The same interaction site as Fig. 8A, embedded in epoxy resin and imaged under confocal microscopy revealed DAB deposited (arrow) beneath and surrounding an attempted penetration site (arrowhead). The deposit was located up against the plant cell wall (PC) on the inside of the cell. Scale bar = 5 μm. doi:10.1371/journal.ppat.1001335.g008
not yet been studied in fungi. A deletion of *tmpL* resulted in a severely reduced virulence defect and hypersensitivity of exogenous oxidative stresses, however when the *YAP1* gene was overexpressed in the deletion line, it rescued these and other mutant phenotypes, suggesting *tmpL*, *YAP1* and presumably *HYR1* may act in a concerted pathway to sense and trigger ROS scavenging pathways.

**MoHYR1 helps the fungus negotiate a hostile host environment**

A successful pathogen, which has the ability to detoxify ROS, will subsequently have fewer barriers to overcome before reaching its ultimate goal, which are the cell contents. Our results with the *MoHYR1* gene suggest that while there might be no effect of *MoHYR1* on ROS that’s produced immediately by the plant (Figure S3), there is subsequent ROS production which *MoHYR1* clearly helps the fungus overcome (Figure 10). Metabolic profiling performed by Talbot and colleagues (2008) provides support for this concept, revealing a *M. oryzae*-induced host metabolism re-programming that suppressed or delayed plant-produced ROS during susceptible interactions.

Although supporting evidence has shown that *M. oryzae* can produce ROS during infection related development [5], through scavenging experiments, the ROS observed in our studies appear to be largely plant-generated. Internal fungal ROS was unaffected by the absence of the *MoHYR1* gene *in vitro*. Furthermore, ROS halos were not disrupted by the ROS scavenger, ascorbic acid, when applied only to conidia, but were disrupted when ascorbic acid was specifically applied to leaves. Several pathways for plant-generated ROS include cell wall-bound peroxidases [1]. Plants defend themselves against pathogens by a battery of cell wall-associated defense reactions, including generation of ROS and cross-linking of lignin compounds [34]. During the interaction between a French bean (*Phaseolus vulgaris*) and a cell wall elicitor from *Colletotrichum lindemuthianum*, ROS appears to originate from cell wall peroxidases [36]. Apoplastic alkalization has been shown to be important in this process [34]. ROS generated from cell wall peroxidases also serve as key molecules required for lignification and cross-linking of cell walls [34]. In a study carried out between barley and the powdery mildew fungus, barley cell wall localized peroxidase *HvRBOHA* is responsible for generating H$_2$O$_2$, which was only present in non-penetrated cells [37]. Our results, particularly in Figure 8B, suggest ROS localized up against the plant cell wall. Further investigations into *M. oryzae*-host interactions will include analyzing plant defense-related genes, including the barley cell wall peroxidase.

Callose and ROS are two plant defensive compounds known to be involved in cell wall appositions, which are deposited during both compatible and incompatible interactions [34]. H$_2$O$_2$ played an important role in this process and enzymatic removal of H$_2$O$_2$ by catalase significantly reduces the frequency of phenolic deposition [34]. Several components were reported to be essential for this oxidative burst: peroxidases, a calcium influx and K$^+$ Cl$^-$ efflux, extracellular alkalization, and post-Golgi vesicles [38]. ROS around the CWA areas might function as signal compounds to gather the vesicles and components needed for mature CWAs. We observed that ROS and callose deposits were positionally related during attempted penetration by both wild type and *Ahp1* mutants, immediately below the appressorium. From this result, we hypothesize that ROS generated by plant defenses activates CWA formation in a susceptible host and experiments to determine the timing of deposition of ROS versus callose are currently underway.

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**Figure 9. ROS scavenging in the plant rescued the hyr1 mutant phenotype.** *(A)* Conidia of *Ahp1* (B25) were mixed with 0.5 mM ascorbic acid and inoculated onto the leaf surface. Infected leaves were stained for ROS 24 hpi. *(B)* Conidia were mixed with water and inoculated onto the leaf surface. Leaves were first treated with 0.5 mM ascorbic acid for 1 hour and then stained for ROS 24 hpi. *(C)* From left to right: *Ahp1* (B25), *Ahp1* (B33), (where susceptible barley leaves were treated with 0.5 mM ascorbic acid for 1 hour and then inoculated with mutant spores in water) ectopic (B40), ectopic (B60), wild type (70-15). Scale bar = 20 μm for all confocal images.

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A hypothesis that follows from these data is that when the *MoHYR1* gene is deleted, the plant responds as though it’s being challenged with an avirulent pathogen. As early as 12 hours post inoculation, we observed that barley leaves inoculated with *Dhyr1* mutants showed higher ROS signals compared with leaves inoculated with wild type. These data were consistent using two staining methods, H$_2$DCFDA and DAB. In leaves inoculated with wild type, ROS was detected around appressoria but was mostly

![Images of plant leaves and appressoria with DAB staining.](image)

**Figure 10. Mutants have more DAB staining than wild type revealed a stronger plant reaction.** DAB staining was performed on wild type (70-15) conidia (A, C, E) and *Dhyr1* (B25) mutant conidia (B, D, F) 24 hpi. Wild type (70-15) conidia on the leaf surface shows DAB staining mostly the fungal structures while *Dhyr1* (B25) mutant conidia elicit a stronger ROS plant reaction. Images were generated with a transmitted light microscope. Scale bars = 100 μm.

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A hypothesis that follows from these data is that when the *MoHYR1* gene is deleted, the plant responds as though it’s being challenged with an avirulent pathogen. As early as 12 hours post inoculation, we observed that barley leaves inoculated with *Dhyr1* mutants showed higher ROS signals compared with leaves inoculated with wild type. These data were consistent using two staining methods, H$_2$DCFDA and DAB. In leaves inoculated with wild type, ROS was detected around appressoria but was mostly

![Images of plant leaves with correlated plant reaction images.](image)

**Figure 11. Two plant defense responses overlap when the *Dhyr1* mutant conidia were inoculated onto leaves.** Correlative images show plant reaction underneath appressoria 24 hpi. (A) ROS staining; (B) aniline blue staining; (C) merged image of panels A and B. Images were processed sequentially (ROS followed by aniline blue), imaged by confocal microscopy and correlated. Scale bar = 2.5 μm.

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observed inside fungal structures. However, ROS was seen both around appressoria and adjacent cells when inoculated with the \( \Delta \text{hyr1} \) mutants. Whole cells filled with ROS were also observed when inoculated with \( \Delta \text{hyr1} \) mutants, which was related with HR-type cell death. All these data indicated that \( \text{HYR1} \) might function to suppress later plant-generated ROS, either by detoxifying it directly, or manipulating plant ROS secretion-related gene expression.

\( \text{MoHYR1} \) regulates several genes involved in ROS-scavenging

While our data showed that \( \text{HYR1} \) likely played an important role in ROS-detoxification processes, our experiments did not preclude other ROS tolerance mechanisms in the fungus, particularly since mutants were reduced in virulence, but not completely non-pathogenic. Such mechanisms might involve the aforementioned \( \text{DESI} \) and \( \text{tup} \) genes. Currently, we are characterizing the \( \text{MoYAP1} \) homolog in \( \text{M. oryzae} \); our initial \( \text{AaAp1} \) mutant data suggested this gene was dispensable for pathogenicity, much like what has been found in \( \text{Botrytis cinerea} \), \( \text{Aspergillus fumigatus} \) and \( \text{Cochliobolus heterostrophus} \) [23,25,26]. Intriguingly, \( \text{YAP1} \) did appear to be essential for virulence in \( \text{Ustilago maydis} \) and \( \text{Alternaria alternata} \) [20,25], suggesting that fungal lifestyle (i.e. necrotrrophic vs. biotrophic) had little to do with this particular oxidative stress pathway, and further supporting redundant pathways. Our real-time qRT-PCR data showed that \( \text{YAP1} \) increases in expression when wild type was challenged with \( \text{H}_2\text{O}_2 \) and we also noted a decrease in \( \text{YAP1} \) gene expression in the \( \Delta \text{hyr1} \) mutant background. One interpretation of this result was that the fungal cell might be compensating for the absence of \( \text{HYR1} \), by boosting expression of its partner gene.

The glutathione pathway-related genes \( \text{GLR1}, \text{GTO1} \) and \( \text{GSH1} \), all increased during \( \text{H}_2\text{O}_2 \) challenge in the wild type however had extremely decreased expression in the mutant line, regardless of ROS. This suggested that these genes were reliant upon \( \text{HYR1} \), which was not unexpected, since the glutathione pathway was shown to be regulated \( \text{YAP1} \), which occurs after interacting with \( \text{HYR1} \) [17]. Our results were also in keeping with the \( \text{C. heterostrophus Yap1} \) homolog mutant \( \text{AaHypp1} \), which showed extremely low levels of both \( \text{GLR1} \) and \( \text{GSH1} \) [23]. Interestingly, we did not observe any of the other genes increasing in expression in the mutant background; this suggested that at least for the genes that we chose such as \( \text{CAT1} \) and \( \text{SOD1} \), they did not provide compensatory mechanisms for a loss of \( \text{HYR1} \). While this is one hypothesis, it is also possible that these genes are regulated at the protein level, as was found in the \( \text{A. fumigatus} \) mutant, \( \text{AaHypp1} \); both \( \text{CAT1} \) and \( \text{SOD1} \) were among the proteins down-regulated in the mutant [39], and this could also hold true for the \( \text{AaYap1} \) mutant. Likewise, catalase, SOD and peroxidase activities were measured in the \( \text{A. alternata} \) mutant \( \text{AaAap1} \) [25]. A transcriptomic study on the \( \text{AaAap1} \) deletion mutant would answer many of these questions; further, such a study would uncover redundant pathways of ROS detoxification masked by the presence of \( \text{MoHYR1} \).

Localization of the \( \text{MoHYR1} \) protein

While numerous studies have examined localization of the \( \text{Yap1p} \), we were unable to find any studies on the localization of \( \text{HYR1} \) either in yeast or filamentous fungi. Our data revealed that the \( \text{HYR1} \) protein mostly localized either to the cytosol or to vacuoles, during early stage infection events on barley (germ tube, early appressorial formation, appressorial maturation and penetration). At one hpi, \( \text{MoHYR1} \) was mainly moving through the germ tube, although it was difficult to definitively ascertain which organelle it might be associated with. At twelve hpi, the \( \text{MoHYR1} \) protein shows cytoplasmic localization, mainly expressed in the cytosol of the appressorium. We suspect that by twenty-four hours, the fungus had penetrated and gained ingress to the first epidermal cell; indeed cell biology studies on events following initial penetration suggested that \( \text{M. oryzae} \) bulbous hyphae fill an entire rice leaf sheath cell and were in the process of moving onto the next one by twenty-seven hours post-inoculation [40]. Its vacuolar localization at this time-point could reflect that fact that it was no longer needed by the fungus, which had circumvented the plant’s oxidative burst and at that point growing in the first epidermal cell. We examined a later time-point at 72 hpi and found the \( \text{HYR1} \),
Figure 13. Antioxidant gene orthologs have altered expression in the \textit{A. thailandica} mutant versus wild type. Wild type (70-15) and \textit{A. thailandica} mutant (B25) were grown in 0 mM and 5 mM hydrogen peroxide and collected 1 hour after immersion. RNA was extracted and real-time qRT-PCR performed on three biological replicates. (A) The \textit{YAP1}, \textit{GTO1}, \textit{GLR1} and \textit{GSH1} all increase in expression in wild type upon \textit{H}_2\textit{O}_2 challenge, but the latter three display low levels in the mutant. (B) \textit{CAT1}, \textit{SOD1}, \textit{GSH2}, \textit{GTL1} and cyt \textit{c} peroxidase do not display significant changes in expression. \textit{MoHYR1} expression is abolished in the mutants. Letters over bars represent statistically significant differences between expression changes of the genes (statistics were generated using student t-test with p-value <0.05).

Conclusions and future directions

In conclusion, we identified and characterized the \textit{MoHYR1} gene, a functional homolog of the yeast \textit{Hyr1} (or \textit{Gpx3}) gene. Although \textit{MoHYR1} does not cause dramatic effects in the disease phenotype, it nevertheless played an important role in virulence. This effect appeared to be related to the deletion mutant’s inability to tolerate plant-generated ROS, or at least to do so in a timely and effective manner to cause wild type levels of disease. Together, our results help to define a mechanism that, while well-studied in yeast, has not yet been examined in filamentous fungi; furthermore, our studies pose additional questions to be answered regarding the role of the glutathione pathway in scavenging ROS in filamentous fungi, how this aids in pathogenicity and what other underlying redundant scavenging pathways exist.

Materials and Methods

\textit{M. oryzae} strains and growth conditions

Rice-infecting \textit{M. oryzae}, strain 70–15 (Fungal Genetics Stock Center 8958) was used as the wild type strain throughout this project, and the strain from which mutants and transgens were derived. All strains were maintained at 25°C under constant fluorescent light on complete medium (CM 1 liter: 10 g sucrose, 6 g yeast extract, 6 g casamino acid, 1 ml trace element). Oatmeal agar medium (OAM 1 liter: 50 g oatmeal and 15 g agar) was used as an oatmeal, and cyt \textit{c} peroxidase do not display significant changes in expression. \textit{MoHYR1} expression is abolished in the mutants. Letters over bars represent statistically significant differences between expression changes of the genes (statistics were generated using student t-test with p-value <0.05).

Yeast strains and complementation assays

Yeast strains BY4741 (wild type) and BY4741 YIR037W (\textit{A. thailandica} mutant) were ordered from the American Type Culture Collection, grown out and maintained on YPD medium. Constructs for transformation were built using standard PCR reaction conditions and programs; briefly, pJS371 used overlapping primers to make an intron-free version of the \textit{MoHYR1} gene in pJS318. Using the intron-free plasmid, overlapping primers were used to make Cys39Ala and Cys88Ala mutant versions of the \textit{MoHYR1} gene.

Figure 14. \textit{MoHYR1} changed localization during pre-penetration events on the surface of a leaf. The \textit{MoHYR1} coding sequence was fused to the cerulean fluorescent protein to study protein localization during early infection. (A) \textit{HYR1} at 1 hpi with putative vacuole location and low level cytoplasmic distribution; the germ tubes have formed, but no appressorium. (B) \textit{HYR1} at 6 hpi with increased cytoplasmic localization where it is likely to be required to function in ROS scavenging; an immature appressorium was apparent. (C) \textit{HYR1} at 12 hpi with cytoplasmic location; a mature appressorium was apparent. (D) \textit{HYR1} at 24 hpi with vacuole and low level cytoplasmic localization in the appressorium. (E) \textit{HYR1} at 72 hpi again showing cytoplasmic localization. Images were taken with confocal microscopy and all experiments were done on the surface of barley leaves. Scale bar shown = 10 μm for all images.

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pJ8382 SchYR1-Pro::MoHYR1_Cys36Ala::SchYR1Term; pJ8383 SchYR1-Pro::MoHYR1_Cys36Ala::SchYR1Term.

**Plants cultivars and growth conditions**

Rice cultivar Maratelli (a gift from the Dean Lab; Raleigh, NC) and barley cultivar Lacey (Johnny’s Selected Seeds; Winslow, ME) were used throughout this study, as both are susceptible to *M. oryzae* strain 70–15. Rice was grown in a growth chamber at 80% humidity, and 12 h:12 h day/night cycles, at 28°C. Barley was grown in a growth chamber at 60% humidity, and 12 h:12 h day/night cycles, at 24°C (day) and 22°C (night).

**Targeted deletion of Hyr1**

The targeted gene deletion was accomplished using the homologous recombination method. We amplified 5′ and 3′ flanking regions of *Hyr1* using primer pairs #1 and 2 (Table S2). Flanking regions were then linked via adaptor-mediated PCR to a 1.3 kb *HPH* coding sequence, providing resistance to the antibiotic hygromycin (Alexis Biochemicals, San Diego, CA). The entire length of the deletion fragment was 3.7 kb. Fungal protoplasts of the wild type 70-15 were directly transformed with the nested PCR product (primers used were forward primer of primer pair #1 and reverse primer of primer pair #2). Protoplast generation and subsequent transformation were conducted by following established procedures [41]. To confirm the knockout mutant, the genomic DNA of candidate strains was extracted and amplified with primer pairs #3, 4 and 5 (Table S2).

**In vitro H₂O₂ growth assessment of Δhyr1 mutants**

Equal-sized pieces of mycelia were cut with a 3 cork-borer tool (0.7 cm in diameter), and immersed in 10 ml of liquid CM at 25°C in darkness. Colonies were grown in CM containing H₂O₂ at concentrations of 0 mM, 5 mM and 10 mM. Colonies were removed from each well, vacuum filtered to dryness, and measured on a scale one week post-inoculation.

**Pathogenicity assays**

For point or drop inoculations, conidia were harvested from 12-day-old cultures grown on OMA in 20 μl of a 0.2% gelatin (Acros organics, New Jersey) suspension, for a final concentration of 1–5×10⁷ conidia/ml. Point two percent gelatin was used as a non-inoculated control for pathogenicity assays. For drop inoculations, three week old leaves of Maratelli or Lacey were detached and laid flat in a humid chamber (90 mm Petri dish with moist filter paper). Twenty microliters of conidial suspensions, or gelatin alone, were dropped onto each leaf and kept in darkness overnight at ~25°C. The next day, remaining water droplets were wicked off and moved to a growth chamber under constant fluorescent light. For spray inoculations, conidial suspensions (10 ml; concentration as above) in 0.2% gelatin were sprayed onto three week old Maratelli or Lacey seedlings. Inoculated plants were placed in a dew chamber at 25°C for 24 hours in the dark, and then transferred into the growth chamber with a photoperiod of 16 h:8 h light:dark cycles. Disease severity was assessed seven days after inoculation.

**Quantitative real-time RT-PCR of ROS-related genes and data processing**

Quantitative real time reverse transcription PCR (real-time qRT-PCR) was carried out using primer pairs for the following genes: YAP1 (MGG_12814.6), GSH1 (MGG_07317.6), GSH2 (MGG_06454.6), GLO1 (MGG_12749.6), GTT1 (MGG_06747.6), GTO1 (MGG_05677.6), GTO2 (MGG_05678.6), SOD1 (MGG_09138.6), CHT1 (MGG_10061.6) and cytochrome c peroxidase (MGG_10368.6). The housekeeping gene encoding ubiquitin conjugating enzyme (MGG_00604.6) was used as an internal control. We also included the gene MoHYR1 (MGG_07460.6) to confirm its deletion in the mutant lines. Primer pairs are listed in Table S3. Seventy-five nanograms of cDNA generated from mycelium grown as per the H₂O₂ experiments described above (generated from the 0 mM and 5 mM H₂O₂ samples), was used as templates for each reaction. The mycelia were fragmented in a blender as per the protocol by Mosquera et al [42], before being inoculated into liquid complete medium. After 2–3 days, the mycelia were blended again to ensure the maximum amount of actively growing fungal tip. The H₂O₂ experiment was performed 24 hours after the 2nd blending, and RNA was extracted. PCR reaction conditions were as follows for a 25 μl reaction: 13 μl H₂O₂, 10 μl 5 Prime SYBR Green Master Mix (Fisher Scientific, Waltham, MA), 0.5 μl Forward Primer (for a final concentration of 2 μM; Integrated DNA Technologies, Coralville, IA), 0.5 μl Reverse Primer (for a final concentration of 2 μM) and 1 μl template DNA. Conditions for real-time quantitative RT-PCR conditions were as follows: 95°C for 2 min; 95°C for 15 sec, 50°C for 15 sec, 68°C for 20 sec (cycle 40 times); 95°C for 15 sec; 60°C for 15 sec (melting curve); 60°C − 95°C for 20 min; 95°C for 15 sec; lid temperature constant at 105°C. The 2−ΔΔCt method was used for generating the data. ΔΔCt is defined as ΔCt treatment - ΔCt calibrator. cDNA from the strain 70-15 in 0 mM H₂O₂ was used as the calibrator for comparison of gene expression in 5 mM H₂O₂ in both the Δhyr1 deletion lines as well as the wild type For both the ΔCt treatment and ΔCt calibrator, ΔCt is defined as Ct gene - Ct housekeeping- gene. For the calibrator, which is 0 μM H₂O₂, this value would be 2⁻¹ or 1. These experiments were repeated twice with similar results.

**Cloning of MoHYR1 and generation of fusion protein**

A Hyr1 N-terminal cerulean fusion construct was generated by fusion PCR. Briefly, using *M. oryzae* genomic DNA as a template, a 1 kb promoter region of *Hyr1* was amplified with primers 6 and 7 (Table S2). Another set of primers, 8 and 9, were used to amplify the 2.4 kb *Hyr1* open reading frame. Three resulting fragments, the 1 kb promoter fragment, the 1328 bp ORF (including 709 bp of terminator sequence) and 740 bp cerulean fluorescent protein coding sequence [43], were mixed and subjected to a second fusion PCR with primers 7 and 8. The resulting 3.1 kb PCR product was generated with *BanIII* and *NotI* restriction enzymes (New England Biolabs, Beverly, MA) and cloned into pBlueScript SK+ pBS. The construct was fully sequenced and found to be correct, hence was co-transformed into the *M. oryzae* Δhyr1 knockout mutant protoplasts to make Cerulean-HYR1 fusion transformants. Transformants with expected genetic integration events were identified by PCR using primers pairs 6 and 10 (Table S2). Properly transformed Δhyr1 mutants were also used as the complemented lines, in Figures 3 and 4, designated as “hyr1-C”.

**Detection of ROS**

Ten-fourteen day old rice and eight day old barley plants were used and collected 24 hours after being inoculated with 10–12 day old conidia (methods as described above). All staining procedures were performed with both rice and barley, however barley was best-suited for microscopy, hence all micrographs shown in this study are of barley. For experiments with 29,79-dichlorofluorescin diacetate (H₂DCFDA) (Invitrogen, Carlsbad, CA), inoculated tissue was collected and incubated for 60 min at room temperature in 5–20 mM H₂DCFDA dissolved in DMSO (less than 0.005% final concentration), then washed with 0.1 mM KCl,
0.1 mM CaCl2 (pH 6.0) and left for 60 min at 22°C before experimentation. Dye excitation was at 488 nm; emitted light was detected with a 500–550 band pass emission filter. DAB staining was carried out using the protocol developed by Thordal Christensen et al [44]. Briefly, leaves were cut at the base with a razor blade and placed in a 1 mg/mL solution of DAB for 0 h under darkness at room temperature. Leaves were decolorized by immersion in ethanol (96%) for 4 h followed by 2 hours in PBS buffer before imaging. A third method of ROS detection was employed for examining ROS internal to, or secreted from, the fungus. Nitroblue tetrazolium (Sigma-Aldrich, St. Louis) was used at 4 mg/mL (in deionized water) and the staining performed for 5 min–30 min at room temperature prior to observation.

**ROS scavenging treatments**

In order to eliminate the ROS generated by fungus, conidia of *Ahyr1* (B25) and wild type (70-15) were mixed with 0.5 mM ascorbic acid (AsA) and inoculated onto the leaf surface. Leaves were stained for ROS at 24 hpi. In order to eliminate ROS generated from the plant, leaves were first treated with 0.5 mM ascorbic acid for 1 hour. To remove excess AsA, leaves were then washed with 0.1 mM KCl, 0.1 mM CaCl2 (pH 6.0) buffer three times for 3 minutes each. Finally, leaves were inoculated with conidia 1hpi and stained for ROS 24 hpi. Additionally, barley leaves were injected with 5 μM DPI (diphenyleneiodonium; Sigma, St Louis), then washed and inoculated, as above.

**Detection of fungal cell wall**

Calcofluor White M2R (Fluorescent brightener 28, F-6258, Sigma, St Louis) was used for detection of the fungal cell wall. We made 10,000-fold dilutions from a saturated Calcofluor White stock solution. For experiments involving conidia on gel-bond (VWR, Arlington Heights, IL), Calcofluor White was applied 1, 4, 8, 12, and 24 hours post inoculation, incubated for 15 minutes, then gently rinsed off with 1X PBS buffer. For experiments involving inoculated plants, inoculated or non-inoculated (control) leaf tissue was collected and immersed in working solution for 15 minutes, then gently rinsed with 0.1 mM KCl, 0.1 mM CaCl2 (pH 6.0).

**Detection of Cell Wall Appositions (CWAs)**

For CWAs staining, we cleared inoculated or non-inoculated (control) leaves in ethanol:acetic acid (6:1 v/v) overnight and washed them with water. Subsequently, cleared leaves were incubated in 0.05% aniline blue (w/v) in 0.067 M K2HPO4 buffer at pH 9.2 overnight and rinsed gently in sterilized deionized water for microscopy.

**Localization of DAB**

Inoculated barley leaves were stained using DAB and rinsed several times in PBS. Thereafter, samples were fixed in 2% paraformaldehyde (Electron Microscopy Sciences, Hatfield, PA) and 2% glutaraldehyde (Electron Microscopy Sciences, Hatfield, PA) in sodium cacodylate (Electron Microscopy Sciences, Hatfield, PA) buffer for 1 hour overnight. Samples were then rinsed three times, 15 min each, in sodium cacodylate and post-fixed with 2% OsO4 in sodium cacodylate for 3–5 hours on a rotator. Again, samples were then rinsed three times for 15 min each, with water on a rotator. Samples then underwent an ethanol dehydration series (25%, 50%, 80% ETOH; 20 min each) on a rotator. Samples were primed with 1% gamma-glycidoxylpropyl trimethoxysilane in 80% ETOH overnight at room temperature and then washed three times for 15 min each in 100% ETOH on a rotator. Samples then underwent a series of infiltrations on a rotator as follows: 100% ETOH/n-BGE (Electron Microscopy Sciences, Hatfield, PA) (1:1) for 30 min, 100% n-BGE for 30 min, n-BGE/Quetol-651 (Electron Microscopy Sciences, Hatfield, PA) (1:3) for 1 hour, n-BGE/Quetol-651 (1:1) for 1 hour, n-BGE/Quetol-651 (3:1) for 1 hour, 100% Quetol-651 for 1 hour, 100% Quetol-651 overnight and 100% Quetol-651 for 1 hour. Finally, samples were embedded and polymerized in an oven at 60°C for about 24 hours.

**Bioinformatic and statistical analyses**

BlastP analysis was done against the fully sequenced genomic database of *M. oryzae* housed at the Broad Institute, using an e-value of 1e-3. ClustalW (X2) was used to perform the full alignment and generate the phylogenetic tree. The final tree image was generated with Tree Viewer. The HYR1 protein secondary structure was predicted using the PSIPRED protein structure prediction server. The structural image of the HYR1 protein was created using the PyMOL molecular viewer. All student t-tests were performed using JMP8 (SAS Institute Inc. 2007. <Title>., Cary, NC: SAS Institute Inc.).

**Confocal microscopy**

Confocal images were taken with Zeiss LSM510 or Zeiss LSM5 DUO using a C-Apochromat 40X NA = 1.2 water immersion objective lens. H2DCFDA ester was excited at 488 nm and fluorescence was detected using a 505–550 nm band pass filter. Calcofluor white was excited at 405 nm and detected using 420–470 nm band pass filter. Cerulean was excited at 438 nm and detected using a 475 long pass filter. We also used transmitted light and reflected light for some confocal experiments.

**Supporting Information**

**Figure S1** Successful deletion of the *HYR1* via homologous recombination of a single insert. (A) Diagram of strategy used for homologous recombination of *HYR1*. The arrow depicts directionality of gene MMG_07460.6, and FS stands for flanking sequence. HygR is the hygromycin phosphotransferase gene (HPH) that confers resistance to organisms that express it. Physical positions of the gene and flanking regions (from supercontig 20) are shown above the diagram. Bottom diagram shows the gene deletion construct that was PCR-ed and linked via adapters. Purple arrows indicate primer sites for determining insertion site (result shown in C). The bottom-most line indicates *HindIII* cut sites for the Southern blot, and positioning of the HPH probe. (B) External flanking region PCR indicates the insert is located in the correct position in the genome (lane loading from left to right: *Ahyr1* B23, *Ahyr1* B33, *Ahyr1* B54, ectopic B40). The size product is the expected ~1.5kb, as based upon the primer positions in A. Gene specific primers indicate that the knockout mutant does not have *HYR1* gene. HPH specific primers indicate the HPH inserted in the genome. (C) Southern blot indicates a single insertion of the construct in the *Ahyr1* mutants. (D) Diagram of the construct used to complement the *Ahyr1* mutant; the cerulean fluorescent protein (CPF) is driven by the native MoHYR1 promoter and linked the N-terminus of the MoHYR1 gene. (E) Southern blot on the complemented mutant line *hyr1-*C probed with the MoHYR1 gene, which revealed four insertions.

Found at: doi:10.1371/journal.ppat.1001335.s001 (1.25 MB TIF)

**Figure S2** *Ahyr1* cannot grow at increased levels of hydrogen peroxide. (A) *Ahyr1* (B23, B33) growth was inhibited at increased levels of hydrogen peroxide (top = 0 mM; middle = 5 mM; bottom = 10 mM) compared to the complemented strain (*hyr1-*C).
C), wild type (70-15) and Ecotypic (B40, B60). (B) Quantification (dry weight) of samples grown in hydrogen peroxide. This experiment was repeated in triplicate with similar results. Different letters over the bars indicate a significant difference as determined by a student’s t-test and a p-value of < 0.05. Found at: doi:10.1371/journal.ppat.1001335.s002 (1.90 MB TIF)

**Figure S3** Ahyr1 accumulated similar levels of ROS to wild type *in vitro*. Hyphae of wild type and *Ahyr1* were grown on complete media plates and stained with nitroblue tetrazolium (NBT) and showed similar staining. A, B, C, and D are microscope images of panels E and F. A, C, and E represent *Ahyr1* (B25) and B, D, F represent wild type (70-15). Scale bars = 100 μm. Found at: doi:10.1371/journal.ppat.1001335.s003 (3.72 MB TIF)

**Figure S4** nox1 and nox2 mutants have same ROS production with wild type on plant 24hpi. A loss of NADPH oxidases in *M. oryzae* does not appear to have a significant effect on ROS haloes. (A-F) Confocal images of the *nox1*, *nox2* and wild type parent lines stained with Calcofluor White (CW) for cell wall visualization and the ROS detector *H₂DCFDA*. The left-most panels show multiple spores and appressorium, while the right-hand panels focus on a representative appressorium (bottom-left: *H₂DCFDA*, bottom-right: CW, top: merge). (G) Graphical representation of the data collected in A showing no significant difference between ROS haloes amongst the strains. Experiments were repeated three times with similar results. Different letters over the bars indicate a significant difference as determined by a student’s t-test and a p-value of < 0.05. Scale bar = 10μm. Found at: doi:10.1371/journal.ppat.1001335.s004 (1.08 MB TIF)

**Figure S5** *Ahyr1* displays similar levels of ROS to wild type immediately after inoculation. (A) ROS signals are detected in barley leaves 1 hpi with either the *Ahyr1* mutants or the wild type strain. *Ahyr1* mutants did not show a defect compared to wild type. Leaves treated with pathogens are significantly brighter than untreated leaves. (B) Quantification of ROS signal intensity reveals a significant difference between inoculated and untreated barley leaves. This experiment was repeated in triplicate with similar results. Different letters over the bars indicate a significant difference as determined by a student’s t-test, and a p-value of < 0.05. Images are taken with confocal microscope. Scale bar = 20 μm. Found at: doi:10.1371/journal.ppat.1001335.s005 (0.77 MB TIF)

**Table S1** *HYR1* amino acid sequence of *M. oryzae* is most closely related to *N. crassa*. Percent identities and similarities were determined using BlastP for ten filamentous fungi, one yeast and one mammal. Found at: doi:10.1371/journal.ppat.1001335.s006 (0.01 MB XLSX)

**Table S2** Primers pairs used to generate the HYR1 deletion construct and to test the targeted deletions. Found at: doi:10.1371/journal.ppat.1001335.s007 (0.01 MB XLSX)

**Table S3** Primers pairs used in real-time qRT-PCR experiments. Found at: doi:10.1371/journal.ppat.1001335.s008 (0.01 MB XLSX)

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**Author Contributions**

Conceived and designed the experiments: KH KJC JLC JAS NMD. Performed the experiments: KH KJC JLC JAS NMD. Analyzed the data: KH KJC JLC JAS NMD. Contributed reagents/materials/analysis tools: KH KJC JLC JAS NMD. Wrote the paper: KH KJC JLC JAS NMD.

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