Antioxidant genes of plants and fungal pathogens are distinctly regulated during disease development in different Rhizoctonia solani pathosystems

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Abstract

Biotic stress, as a result of plant-pathogen interactions, induces the accumulation of reactive oxygen species in the cells, causing severe oxidative damage to plants and pathogens. To overcome this damage, both the host and pathogen have developed antioxidant systems to quench excess ROS and keep ROS production and scavenging systems under control. Data on ROS-scavenging systems in the necrotrophic plant pathogen Rhizoctonia solani are just emerging. We formerly identified vitamin B6 biosynthetic machinery of R. solani AG3 as a powerful antioxidant exhibiting a high ability to quench ROS, similar to CATALASE (CAT) and GLUTATHIONE S-TRANSFERASE (GST). Here, we provide evidence on the involvement of R. solani vitamin B6 biosynthetic pathway genes; RsolPDX1 (KF620111.1), RsolPDX2 (KF620112.1), and RsolPLR (KJ395592.1) in vitamin B6 de novo biosynthesis by yeast complementation assays. Since gene expression studies focusing on oxidative stress responses of both the plant and the pathogen following R. solani infection are very limited, this study is the first coexpression analysis of genes encoding vitamin B6, CAT and GST in plant and fungal tissues of three pathosystems during interaction of different AG groups of R. solani with their respective hosts. The findings indicate that distinct expression patterns of fungal and host antioxidant genes were correlated in necrotic tissues and their surrounding areas in each of the three R. solani pathosystems: potato sprout-R. solani AG3; soybean hypocotyl-R. solani AG4 and soybean leaves-R. solani AG1-IA interactions. Levels of ROS increased in all types of potato and soybean tissues, and in fungal hyphae following infection of R. solani AGs as determined by non-fluorescence and fluorescence methods using H$_2$DCF-DA and DAB, respectively. Overall, we demonstrate that the co-expression and accumulation of certain plant and pathogen ROS-antioxidant related genes in each pathosystem are highlighted and might be critical during disease development from the plant’s point of view, and in pathogenicity and developing of infection structures from the fungal point of view.
Introduction
The necrotrophic fungus *Rhizoctonia solani* Kühn (teleomorph *Thanatephorus cucumeris*, Frank, Donk) is an economically devastating plant pathogen with a wide host range. *R. solani* is classified into fourteen anastomosis groups (AGs) based on hyphal fusion [1], and strains belonging to AG3 and AG4 are root infecting pathogens that cause damping-off and stem rot of potato (*Solanum tuberosum* L.) and soybean (*Glycine max* (L.) seedlings, respectively [2–5]. Isolates of the *R. solani* AG1-IA complex can infect aerial portions of the plant as in the case of soybean leaves causing rhizoctonia foliar blight (RFB) [3, 6, 7]. The common elements in disease development of *R. solani* isolates are the close association of fungal hyphae with the host epidermis forming branches known as infection cushions or aggregates, penetration of the epidermis, inter- and intracellular colonization and breakage of plant tissue by the production of hydrolytic enzymes, which eventually leads to the development of browning and necrosis associated with oxidative burst and death of tissue [4, 8]. Current cultural and chemical controls are not completely effective to manage Rhizoctonia diseases and the diseases remain a persistent problem. Furthermore, resistance to *R. solani* in any plant species does not exist.

One of the earliest plant responses following pathogen recognition is the hypersensitive response, leading to the production of reactive oxygen species, primarily superoxide (O$_2^-$) and H$_2$O$_2$, at the site of attempted invasion [9]. The produced ROS activates plant defense responses, including programmed cell death, or functions as second messengers in the induction of various plant defense-related genes [10, 11]. In the case of necrotrophic fungi, ROS plays a central role during their interaction with their plant hosts by stimulating the plant’s basal defense responses [12–15]. Several studies showed that the onset of basal resistance in plants to *R. solani* is tied with ROS-scavenging mechanisms, accumulation of metabolites related to vitamin B6 biosynthetic pathway, oxylipins production and cell wall bound phenolic compounds [16–21].

Different strategies of oxidative stress response systems are deployed by biotrophic and necrotrophic fungi during the infection process. Biotrophic pathogens such as rust fungi respond to oxidative stress by containing and suppressing the oxidative burst, while necrotrophic pathogens such as *Botrytis cinerea* rely on the exploitation of the oxidative burst in plants to its own advantage and in some cases contribute to it [22]. Hence, the ability of necrotrophic fungi to surpass or manipulate ROS-related plant defenses is crucial for disease progression, and ROS detoxification is essential to the sensitivity of the necrotrophic fungus while encountering its host. Detoxification systems, such as the NOX complex is a good example of how necrotrophic fungi such as *B. cinerea* and *Alternaria alternata* can sustain a reduced redox states within subcellular microenvironments [22]. In recent years, vitamin B6 was recognized as a strong antioxidant displaying a great capacity to quench ROS matching that of tocopherols or ascorbic acid, and may have a function in stress alleviation in fungi and plants [23, 24]. Data on ROS-scavenging systems in *R. solani* are just emerging. To date, the upregulation of *R. solani* genes particularly, PYRIDOXAL REDUCTASE (PLR AKR8; DW520695), and PYRIDOXAL-5-PHOSPHATASES and TRANSAMINASES of the vitamin B6 salvage biosynthetic pathway, was reported in *R. solani* hyphae in association with a mycoparasite or an antagonistic bacteria, respectively [25, 26].

In our previous publication, we fully characterized two genes of the *de novo* vitamin B6 biosynthetic pathway; *RsolPDX1* (KF620111.1) and *RsolPDX2* (KF620112.1) genes, and one gene *RsolPLR* (KJ395592.1) of the vitamin B6 salvage biosynthetic pathway of *R. solani* AG3. Upon exposure to the ROS-generating chemicals, paraquat and H$_2$O$_2$, the vitamin B6 genes exhibited differential regulation and their transcript abundance levels were mostly higher than levels to the well-recognized antioxidant genes, CATALASE (CAT) and GLUTATHIONE S-TRANSFERASE.
Gene expression studies focusing on oxidative stress responses from both the plant and the pathogen sides during plant- \textit{R. solani} interactions are very limited. Foley, Kidd [28] identified a number of wheat-derived and \textit{R. solani} AG8-derived genes involved in ROS production and redox regulation, and whose expression was affected during infection. Their study highlighted the need to understand the ROS-scavenging system interplay of fungal and host derived genes involved in ROS/redox regulation at the site of infection and the surrounding areas. However our knowledge on the role of vitamin B6 genes as antioxidants in other AGs of \textit{R. solani} is limited.

In this study, we first provided indirect evidence on the functionality of \textit{RsolPDX1} and \textit{RsolPDX2} of \textit{R. solani} AG3, and their involvement in vitamin B6 \textit{de novo} biosynthesis pathway via heterologous complementation of the yeast, \textit{Saccharomyces cerevisiae} strains \textit{Δsnz1} and \textit{Δsno1}. Second, we showed that the antioxidant genes encoding vitamin B6 (i.e., \textit{PDX}, \textit{PLR}), \textit{CAT} and, \textit{GST} of \textit{R. solani} AG3 and potato are differentially induced and transcriptionally regulated at the site of infection (i.e., necrotic tissues, and in the surrounding areas) during \textit{R. solani} AG3-potato sprout interaction (pathosystem I). Third, we extended our study to investigate whether differential and spatial expression patterns and transcriptional regulation of vitamin B6 genes and other antioxidant genes also occur in the pathogen and host upon \textit{R. solani} AG4 infection of soybean hypocotyl (pathosystem II) and also upon \textit{R. solani} AG1-IA infection of soybean leaves (pathosystem III).

**Materials and methods**

**Yeast complementation assays- heterologous expression of \textit{R. solani} AG3 \textit{PDX1} and \textit{PDX2} and growth assays**

Full-length clones of \textit{RsolPDX1-AG3} and \textit{RsolPDX2-AG3} were obtained by primer pairs (\textit{RsolAG3-PDX1-CDS-F/R} and \textit{RsolAG3-PDX2-CDS-F/R}) (Table 1) from cDNA of \textit{R. solani} AG3 isolate Rs114 (ATCC 10183). The PCR products of the two genes were purified using the QIAquick PCR Purification Kit (Qiagen, Toronto, ON, Canada), and sub-cloned into the pDrive cloning vector (Qiagen) to obtain pDrive-\textit{PDX1}-AG3 and pDrive-\textit{PDX2}-AG3. The clones were digested with NotI and ligated into a NotI-digested yeast shuttle vector pFL61 (Addgene, Cambridge, MA, USA) [29] to obtain pFL61-\textit{PDX1-AG3} and pFL61-\textit{PDX2-AG3}. All clones were sequenced to confirm the presence of the target genes. To determine whether \textit{PDX1-AG3} and \textit{PDX2-AG3} encode functional proteins, the \textit{Saccharomyces cerevisiae} mutant strains (\textit{Δsnz1}: disrupted \textit{pdx1} and \textit{Δsno1}: disrupted \textit{pdx2}), obtained from the European \textit{Saccharomyces cerevisiae} Archive for functional analysis (EUROSCARF; http://wwwrz.uni-frankfurt.de/FB/fb16/mikro/EUROSCARF/), were transformed using a lithium acetate-based method [30] with the construct pFL61-expression vector or with pFL61 alone (negative control), and tested for their capacity to complement the defect function of the \textit{snz1} and \textit{sno1}.

For vitamin B6 growth-dependence assays: Yeast cultures grown to exponential phase (2 × 10^7 cells mL^{-1}) followed by two serial 10-fold dilutions corresponding to the wild-type BY4742, and transformants were spotted on minimal selective SC medium (Difco Laboratories, MI, USA) medium lacking vitamin B6. The mutants, \textit{Δsnz1} and \textit{Δsno1} are sensitive to the superoxide generator menadione, and this sensitivity is corrected by the addition of vitamin B6 [31]. Transformed yeast cells were grown on minimal SC medium to the post-diauxic phase for 20h followed by a treatment with 40 mM menadione (-Aldrich Canada, Mississauga, ON, Canada) for 3 h in the presence or absence of 2μg mL^{-1} of vitamin B6 and spotted on SC medium lacking Uracil and vitamin B6. Pictures were captured from plates that were incubated for 3 d at 30°C [32].
Table 1. List of primers used in this study.

| Target organism | Primer | Sequence (5' → 3') | Annealing temperature (°C) | Amplicon size (bp) | Reference |
|-----------------|--------|--------------------|-----------------------------|--------------------|-----------|
| **PDX1 (Pyridoxine biosynthesis gene)** | Soybean | GM-1-PDX1.1-F | TACAGTTATACGGCAACGGGTGCA | 59 | 235 | XM_003542937.3 |
| | | GM-PDX1.1-R | GTCCATGACGATCGGGG | | | |
| | | GM-PDX1.2-F | TACAGCTCGCGCCAAACGGG | 54 | 235 | NM_001251112.2 |
| | | GM-PDX1.2-R | ATCCCTGACGATGGG | | | |
| **Potato** | ST-PDX1.1-qF | CTGCTGACTATCGTGGTAAT | 55 | 83 | [37] |
| | | ST-PDX1.2-qF | TGCTCTATCCCTTACAG | 55 | 162 | |
| **R. solani AG3** | 5'GSP1-(PDX1) | TGACGAACTCAGCGAACAATC | 55 | 219 | [27] |
| | 3'GSP2-(PDX1) | TCCGTTGCTGAGGATCGCTC | | | |
| **R. solani AG4** | ATGAAGCGAACGATCGCTGGA | 53 | 243 | AVOZ01001318.1 |
| | RsolAG8-PDX1-R | GATCAATGCGATCGCTGGA | | | |
| **R. solani AG1-IA** | RsolAG1-IA-PDX1-F | ATCTCTGCAATCGGATTGAG | 53 | 207 | JATN0100314.1 |
| | RsolAG1-IA-PDX1-R | TGCCCTGACGATCGCTGGA | | | |
| **PDX2 (Pyridoxine biosynthesis gene)** | Soybean | GM-PDX2.1-F | TTAGGAGCTGAAAGGTGGTGGAG | 53 | 328 | XM_014774652.1 |
| | | GM-PDX2.1-R | TTGGGAGCAAGAGCTGTCGAC | | | |
| **Potato** | ST-PDX2-qF | ATTCCACATCTCTATTC | 55 | 185 | [37] |
| | ST-PDX2-qR | CACAATACAGGATCGCTC | | | |
| **R. solani AG3** | 5'GSP1-(PDX2) | AATCTTCATGCTGACGATCGCTGGA | 53 | 207 | JATN0100314.1 |
| | 3'GSP2-(PDX2) | TGGCTGCTGAGGATCGCTGGA | | | |
| **R. solani AG4 & R. solani AG1-IA** | RsolAG8-PDX2-F | ATATATACAGCGGATCGCTGGA | 53 | 207 | JATN0100314.1 |
| | RsolAG8-PDX2-R | GGGCCCTTACACCCACACAG | | | |
| **PLR (PYRIDOXYL REDUCTASE)** | Soybean | GM-PLR-F | TCCTCCAAGGCTGAAACGGCA | 53 | 115 | XM_003527250.3 |
| | GM-PLR-R | TAGAAGGAGCTGAGTGGTG | | | |
| **Potato** | ST-PLR-F | GGCAGCCAAAAACGGGACCCGA | 53 | 115 | XM_003527250.3 |
| | ST-PLR-R | AGAAGGAGCTGAGTGGTG | | | |
| **R. solani AG3** | PLR ACR8-F | GAAAGCGTCTCTTGGAAATTC | 53 | 300 | [25] |
| | PLR ACR8-R | GGTTAGATTAGAGTGGCGTGG | | | |
| **R. solani AG4 & R. solani AG1-IA** | RsolAG1-IA-PLR-F | AAGGGGCCAATGCCGAGAAG | 53 | 194 | AFRT01000983.1 |
| | RsolAG1-IA-PLR-R | ATGTTGGCCGAAGGAGGAGA | | | |
| **CATALASE** | Soybean | GM-CAT.4-F | ACATGCTCTCTTTTATTTG | 47 | 149 | NM_001253092 |
| | GM-CAT-R | CTCTATACATTCTCTGTCTT | | | |
| **Potato** | ST-CAT-F | ATGTTGCTCTCTCTCCTGAG | 52 | 147 | AY422179.2 |
| | ST-CAT-R | CTGACACCCACATGCGGGCTT | | | |
| **R. solani AG3** | RsolAG3-CAT-F | ACCAGAAGATTTTAGTCCACGCG | 56 | 190 | [27] |
| | RsolAG3-CAT-R | CATCCCTGTCACAGGCGTGA | | | |
| **R. solani AG4 & R. solani AG1-IA** | RsolAG1-IA-CAT-F | TGGGGGACATGCAGTTGCCCTCTC | 55–54 | 114 | AFRT01002805.1 |
| | RsolAG1-IA-CAT-R | TCCGTTGCTGCTCTCTCCTATTC | | | |
| **GLUTATHIONE S-TRANSFERASES** | Soybean | GM-GST.4-F | TCCATTGGAGAGGTGAGATGC | 53 | 208 | NM_001251762 |
| | GM-GST-R | ATTTCTGACTCTGACCGCCTTAC | | | |

(Continued)
Biological material and inoculation

**Fungal material.** Starter cultures of *R. solani* AG3 isolate Rs114 (ATCC 10183), AG4 isolate A76 obtained from M. Cubeta, North Carolina University, USA, and AG1-IA strain ROS2A4 obtained from P. Ceresini, University of São Paulo State (UNESP), Brazil were recovered from cryogenic storage at −80˚C on potato dextrose agar (PDA; Difco Laboratories) and grown for 3 days at 24˚C in the dark and used as a source of inoculum for plant infection. Cultures of *R. solani* strains grown on half-strength PDA overlaid with cellophane membrane were used for the visualization of ROS in their hyphae.

**Plant material.** Certified potato tubers (*Solanum tuberosum* var. Russet Burbank, susceptible to *R. solani* AG3), obtained from the Potato Research Center (NB, Canada) and propagated at the Macdonald Campus Farm (Ste. Anne de Bellevue, Canada), were kindly provided by D.J. Donnelly (McGill University). Potato tubers were grown without light at 4˚C to induce

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**Table 1. (Continued)**

| Target organism | Primer | Sequence (5’ → 3’) | Annealing temperature (˚C) | Amplicon size (bp) | Reference |
|-----------------|--------|--------------------|---------------------------|-------------------|-----------|
| **Potato**      | ST-GST-F | TCCCTTTAAGCCATAGAGTTGA | 51 | 200 | POTPR1A |
|                 | ST-GST-R | CTTCAAATGCTCTCATCAGTC |                  |                  |           |
| **R. solani AG3** | RsolAG3-GST-F | AGAAGACGAGGCAATAGCAGA | 57 | 256 | [27] |
|                 | RsolAG3-GST-R | ATCTCTTCAAGGGCTTCCAGT |                  |                  |           |
| **R. solani AG4 & R. solani AG1-IA** | RsolAG1-IA-GST-F | TACAACGCATATCCTATCGC | 53 | 217 | ELU40094.1 |
|                 | RsolAG1-IA-GST-R | TTGAGCCGGTAAGTTTTG |                  |                  |           |
| **Reference genes** |        |                    |                           |                  |           |
| **Soybean**     | UKN2/F  | GCCTCTGGATACCTGCTCAAG | 58 | 79  | [38] |
|                 | UKN2/R  | ACCCTCCTCCTAAACTCTCTG |                  |                  |           |
| **Potato**      | ST-Actin7-qF | GGCTATGTATGTTGCTAT | 55 | 186 | [37] |
|                 | ST-Actin7-qR | ATCTTCACTAGTTAAGCAG |                  |                  |           |
| **R. solani AG3** | G3PDH-F | GGATATTATTGATACACTGGA | 55 | 129 | [25] |
|                 | G3PDH-R | TTAAGGCTCAGTCTTTCCT |                  |                  |           |
| **R. solani AG1-IA** | RsolH3-F | CTCCAAATCATCGGCAGTCCCTC | 52 | 76  | [34] |
|                 | RsolH3-R | ATGGTATCTTCTCGAACAAGACAGAG |                  |                  |           |
| **R. solani AG4** | GM-RS-4F | CGTTTCATCTCGATTTACCCT | 55 | 88  | [39] |
|                 | GM-RS-3R | AGTGTATAGTGGTTACCT |                  |                  |           |
| **Transformation** |        |                    |                           |                  |           |
| **R. solani AG3** | RsolAG3-PDX1-CDS-F | GCGGCCGCATCATGTC TGCTCCTG TGTC | 55 | 969 | KF620111.1 |
|                 | RsolAG3-PDX1-CDS-R | GCGGCCGCCTAGCTAGTCTGCT |                  |                  |           |
| **R. solani AG3** | RsolAG3-PDX2-CDS-F | GCGGCCGCATCTCGATCTGCT | 55 | 747 | KF620112.1 |
|                 | RsolAG3-PDX2-CDS-R | GCGGCCGCCTAGCTAGTCTGCT |                  |                  |           |

a Glycine max
b Solanum tuberosum
c Rhizoctonia solani
d Catalase
e Glutathione S-Transferase
f Hypothetical protein unknown 2
g G3PDH: Glyceraldehyde-3-phosphatedehydrogenase
h Histone 3
i ITS1 of 5.85 rRNA

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sprout formation. For sprout induction, uniform-sized tubers were surface sterilised and sprouted for 12 days under controlled conditions following the methods of Aliferis and Jabaji [17], Chamoun, Samsatly [33]. Prevention of browning of the sprout tips was achieved by applying 1.5 mL solution of Ca(NO₃)₂ (0.5M) to sprouts of 3 cm in length. Inoculation of sprouts was performed when sprouts were 8 cm in length.

Soybean (Glycine max) cv. Williams 82 seeds, a susceptible cultivar to R. solani AG-4, were sterilized according to Aliferis, Faubert [16]. Pre-germinated seeds were planted in Cone-tainers® (Stuewe & Sons, Inc., Oregon, USA) filled with 130 mL of sterile turf:perlite (1:1, v/v) and incubated in the dark in a growth chamber in order to produce etiolated seedlings with longer lengths of hypocotyls (approximately 15 cm in height). Hypocotyls were inoculated with AG4 isolate A76 as previously published [16].

For unifoliate leaf inoculation with R. solani AG1-IA, pre-germinated soybean seeds in Cone-tainers® as described above were grown under controlled conditions of temperature, humidity and light following the method of Copley, Aliferis [34]. Fully expanded soybean unifoliate leaves were detached from 2 week-old plants, and used for inoculation.

**Inoculation of potato sprouts, soybean hypocotyls and soybean leaves**

Excised sprouts and etiolated hypocotyls were arranged horizontally in sterile Pyrex trays (40 cm × 26 cm) lined with wet sterile Whatman No. 1 paper. Each treatment replicate consisted of 4 excised potato sprouts or 4 etiolated hypocotyls. There were 3 biological replicates per treatment. Inoculation was performed by sandwiching horizontally the sprouts (8 cm in length) or hypocotyls (15 cm in length) of each treatment replicate between two PDA strips (2 cm × 8 cm) of a 3-day-old R. solani AG3 Rs114 or R. solani AG4 isolate A76 cultures, respectively, with the bottom edge of the strips placed at 4 cm from the basal part of the potato sprouts or hypocotyl sprouts according to the method of Aliferis and Jabaji [17], Chamoun, Samsatly [33]. Potato sprouts or soybean hypocotyls sandwiched with sterile PDA strips served as controls. Moist, sterile absorbent cotton wool was placed at the base of the sprouts or the hypocotyls of each replicate treatment to maintain humidity. Pyrex trays were sealed with Saran Wrap, and placed in a growth chamber under photo-synthetically inactive black light (365 nm) at 24°C.

Fully expanded unifoliate soybean leaves were placed into sterile Pyrex trays as described above. Inoculation was performed by placing PDA plugs (5 mm) of 3-day-old R. solani AG1-IA culture disc, or with sterile PDA alone (control) at the center of leaves. Each replicate treatment consisted of 5 unifoliate leaves. Trays were wrapped with plastic membrane, placed in growth chambers under conditions of day/night temperatures, light cycles and humidity as we previously described in Copley, Aliferis [34].

PDA strips or plugs were removed 120 h, 36h and 18h post-inoculation (HPI) of plant tissue to reveal visible necrotic lesions with infection zones on sprouts, hypocotyls and leaves, respectively. These time points were selected in order to capture the onset of infection cushions and the development of mycelial aggregates that were stereoscopically confirmed (Fig 1). Necrotic tissues containing infection zones plus a one-cm area away from necrotic tissues were harvested. Similar areas and amounts of tissues were harvested from control samples. Harvested tissues were flash-frozen with liquid nitrogen and stored at −80°C. Each treatment (infected or control) consisted of three biological replications.

**RNA extraction, primer design and quantitative RT-PCR**

Total RNA was isolated from 100 mg of flash frozen pulverized tissue obtained from the necrotic lesions and their surrounding areas of respective treatments and corresponding areas of control soybean hypocotyls, leaves, or potato sprouts. Extraction was done using the...
TRIZOL reagent (Generay Biotech, Shanghai, China) following the manufacturer's instructions. RNA concentration and purity was spectrophotometry measured using ND1000 (NanoDrop, Wilmington, Delaware), and RNA quality was verified by gel electrophoresis. RNA (500 ng) was reverse transcribed using the Quantitect Reverse transcriptase kit (Qiagen). QRT-PCR assays were conducted on fungal and plant target antioxidant genes encoding GST, CAT, the de novo vitamin B6 biosynthesis genes; a synthase (PDX1) and a glutaminase, (PDX2), and the vitamin B6 salvage pathway encoding gene, PYRIDOXAL REDUCTASE (PLR) and appropriate internal reference genes for the three pathosystems using Stratagene Mx3000 (Stratagene, Cedar Creek, USA). Primer names were preceded by the plant or fungus abbreviation: GM, Glycine max, ST, Solanum tuberosum, or Rsol, R. solani. Primer sets were designed based on sequences from NCBI, and were checked for specificity to amplify only their target gene. QRT-PCR conditions were optimized for each primer set, and products were confirmed by sequencing. For primer pairs that were used on more than one AG, they were checked for their ability to amplify with comparable efficiencies in the different pathosystems. Reverse transcription PCR assays were performed on three biological replicates and two technical replicates. PCR assay conditions were performed as previously described [27, 35] using suitable annealing temperature for each primer pair (Table 1). In all QRT-PCR assays, routine negative and positive controls were performed at every run. No template control served as negative control. Also, cDNA of uninfected plant host or R. solani alone served as negative controls when amplifying fungal genes or plant genes, respectively. Positive controls for each run consisted of cDNA of R. solani alone or plant host alone when amplifying the fungal or plant genes, respectively. The relative transcript abundance levels of the plant and fungal-derived genes were estimated and normalized against their respective reference genes according to Zhao and Fernald [36]. In the case of R. solani AG-3, expression of the antioxidant genes was normalized using G3PDH. This choice was based on the lowest coefficient variation when compared to Tubulin and Histone using the statistical tool Bestkeeper (http://www.gene-quantification.info).

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Statistical analyses

Significance of the relative transcript abundance between treatments and controls were analyzed by two-way analysis of variance (ANOVA), and when necessary by least significant differences (LSD) at $P < 0.05$ using the SPSS statistical package v. 22.0, (IBM Corp., Armonk, NY, USA). Transcript changes were deemed statistically and biologically significant if $P < 0.05$ and fold changes were $> +1.5$ or $> -1.5$.

In order to find potential correlations and trends between relative abundance of plant and fungal genes expressed in necrotic tissues and their surrounding areas of each pathosystem, a multivariate analysis of the data was performed using the SIMCA-P+ v.12.0 software (Umetrics, MKS Instruments Inc., Andover, MA, USA). The data matrix consisted of biological replicates of control plant tissues, and tissues of necrotic lesions and areas surrounding them (columns; X variables) and the variables for relative transcript abundance of antioxidant genes (rows; Y variables). Data were obtained from the analysis of relative transcript abundance of 11 genes for potato sprouts-\textit{R. solani} AG3, soybean leaves-\textit{R. solani} AG1-IA and soybean hypocotyls-\textit{R. solani} AG4 [11 rows x 9 columns]. For the evaluation of data and detection of outliers, principal component analysis (PCA) was performed. To determine which of the genes were most affected in each pathosystem, PCA loading coefficient plot for the effect of the relative transcript abundance of genes on tissue type, and a loading biplot was built for the visualization of correlations between the X and Y variables using the pc (corr) (i.e., correlation scaled loadings of the correlation between Y variables and X scores based on the variable importance of X as scores and loadings) setting in SIMCA-P+. For data normalization, mean-centering and Pareto [PAR] scaling were used.

Optical and fluorescence microscopy

Association of the detected ROS that accumulated during disease development, with changes in the transcript abundance of the genes was visualized during disease development in the three pathosystems. Cellular and extracellular ROS accumulation was visualized by $2',7'$-dichlorodihydrofluorescein diacetate (H$_2$DCF-DA) and $3,3'$-diaminobenzidine (DAB) staining methods in control and infected soybean leaves and hypocotyls, and potato sprouts, along with mycelia of control \textit{R. solani} AG1-IA, AG4, AG3 grown on half-strength PDA overlaid with cellophane membrane. For ROS detection, samples were incubated with 10 $\mu$M H$_2$DCF-DA, a specific ROS molecular-detection probe, in H$_2$O for 30 min. Samples were then washed with pre-warmed (28˚C) H$_2$O for 30 min to remove the non-internalized probe. To visualize H$_2$O$_2$ accumulation \textit{in situ}, DAB staining was performed, by treating the samples DAB as described in Pogany, von Rad [40] with some modifications. Whole leaves of soybean were treated with DAB for 24h, whereas thin shavings (approximately 2 x 0.5 cm) of soybean hypocotyls and potato sprouts were obtained with a blade and treated for 4h. All samples were cleared with saturated 15.1 M of chloral hydrate solution, and examined under Zeiss SteREO Discovery.V20 microscope (Carl Zeiss Canada Ltd., Toronto, Ontario, Canada). Fluorescence detection from plant and fungal tissues were read at an excitation wavelength of 470 nm using a GFP filter.

Results

Functional characterization of \textit{RsolPDX1} and \textit{RsolPDX2} by yeast heterologous complementation provided proof of the ability of \textit{R. solani} AG3 \textit{PDX1} and \textit{PDX2} to encode functional enzymes

To determine whether \textit{RsolPDX1} and \textit{RsolPDX2} encode functional enzymes, the full coding sequence of each gene was cloned into the yeast expression vector pFL61 and transformed into
S. cerevisiae strains defective in either snz1 (yeast functional PDX1 homolog; Δsnz1) or sno1 (yeast functional PDX2 homolog; Δsno1) and therefore unable to grow in media lacking vitamin B6 [31, 32]. Thus, growth of the transformed mutant S. cerevisiae cells on media lacking vitamin B6 will confirm the complementarity of the R. solani homologs to yeast vitamin B6 genes in addition to their role in vitamin B6 biosynthesis. On media not amended with pyridoxine, growth of S. cerevisiae strains Δsnz1 and Δsno1 expressing RsolPDX1 and RsolPDX2, was evident (Fig 2A and Fig 2B). In contrast, the non-transformed strains or the empty vector-transformed Δsnz1 or Δsno1 mutant cells did not grow on media lacking pyridoxine. Complementation of the Δsnz1 mutant by Rsol-PDX1 and Δsno1 mutant by Rsol-PDX2 demonstrated that Rsol-PDX1 and Rsol-PDX2 are functional, and that they can replace the yeast SNZ1 and SNO1, respectively, implying that they are involved in de novo vitamin B6 biosynthesis.

Δsnz1 and Δsno1 display sensitivity to the superoxide generator menadione, and this sensitivity is corrected by the addition of vitamin B6 [31]. Thus, it is expected that vectors harbouring Rsol-PDX1 or Rsol-PDX2, but not the mutants transformed with the empty vector, are resistant to menadione (Fig 2C and Fig 2D). Unsurprisingly, growth defect restoration of the empty vector-transformed mutants (pFL61-Δsnz1, pFL61-Δsno1) was achieved by the addition

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Fig 2. Complementation of the yeast Δsnz1 and Δsno1 mutants by Rsol-PDX1 and Rsol-PDX2, respectively. Yeast cultures grown to exponential-phase followed by two serial 10-fold dilutions were placed onto vitamin B6-deficient SC (SC-B6) agar plates composed of (A) wild-type (WT), Δsnz1 mutant, Δsnz1 transformed with either pFL61 alone or the RsolPDX1 expression vector (pFL61-RsolPDX1), (B) wild-type (WT), Δsno1 mutant, Δsno1 transformed with either pFL61 alone or the RsolPDX2 expression vector (pFL61-RsolPDX2) (left colonies: 10⁻¹ ml⁻¹ right colonies: 10⁻² ml⁻¹), (C) Δsnz1 mutant yeast transformed either with pFL61 alone or with the RsolPDX1 expression vector (pFL61-RsolPDX1), and (D) Δsno1 mutant yeast transformed either with pFL61 alone or with the RsolPDX2 expression vector (pFL61-RsolPDX2) were both grown to the post-diauxic phase in SC-Uracil medium followed by a 180 min treatment with 40 mM menadione in the presence (+B6) or absence (-B6) of 2 μg ml⁻¹ of vitamin B6. Following menadione treatment, two serial 10-fold dilutions using SC-Uracil medium (left colonies: 10⁻¹ ml⁻¹, right colonies: 10⁻² ml⁻¹) were placed in SC-Uracil-B6 plates. Pictures were obtained after 3 d incubation at 30˚C.
of 2 μg ml⁻¹ of pyridoxine. In conclusion, these data indicate that *Rsol-PDX1* and *Rsol-PDX2* confers resistance to ROS to the cells.

**Distinct expression of host and pathogen antioxidant genes in necrotic tissues and surrounding areas**

For the discovery of trends of plant and fungal antioxidant encoding genes, in the three pathosystems, PCA revealed tight groups with no outliers (*P* < 0.05) (S1 Fig). Antioxidant genes that influenced the separation of treatments (i.e., plant control, infected plant necrotic tissues and surrounding tissues) for each pathosystem were selected based on strong loading coefficients obtained from the loadings plots (Fig 3) and designated to have an effective antioxidant role in plant-pathogen interactions of each pathosystem. The strength of the correlations was pathosystem-dependent. Additionally, selected genes were highly associated with the distinct plant tissue type as revealed by the PCA-loading biplots (Fig 3).

*R. solani* AG3 infection of potato sprouts significantly activates the fungal *RsolAG3-CAT* and *RsolAG3-GST* in addition to the *de novo* vitamin B6 genes of both the pathogen and the host

A remarkable gene upregulation of the fungal and plant vitamin B6 *de novo* pathway genes was observed during disease development of *R. solani* AG3 on potato sprouts. Based on PCA loading biplots and coefficients, the model showed that the loading coefficient values (p) of *RsolAG3-PDX1* (p, 54%) ST-*PDX1.1* (p, 40%), *RsolAG3-PDX2* (p, 35%), and *RsolAG3-GST* (p, 25%), were tightly linked with tissues of surrounding areas of necrotic lesions. On the other hand, the potato *CATALASE*, *ST-CAT* (p, -45%) was negatively associated with necrotic tissues and their surrounding areas (Fig 3A and Fig 3B). Significant fold increases of transcript abundances of the above fungal genes ranged from a minimum of 1.6 (*P* = 0.0029) to a maximum of 4.7 (*P* < 0.0001) depending on the gene and the plant’s area sampled (i.e., necrotic tissue vs surrounding or adjacent tissue) for *RsolAG3-PDX1* and *RsolAG3-PDX2* (Fig 4). Interestingly, the plant homologue ST-*PDX1.1* was differentially upregulated and tissue-dependent (Fig 4). ST-*PDX1.1* had a significant increase (1.9 folds; *P* = 0.0006) in only the tissues surrounding the necrotic area. Significant activation of fungal *RsolAG3-CAT* (70.9 folds; *P* < 0.0001) and *RsolAG3-GST* (12.4 folds; *P* < 0.0001) was observed in necrotic tissues and surrounding areas, respectively.

Fungal-derived vitamin B6 genes and *RsolAG4-GST* are differentially regulated during soybean root and stem rot disease development caused by *R. solani* AG4

The fungal vitamin B6 genes along with *RsolAG4-GST* play a significant role in soybean-*R. solani* AG4 pathosystem compared to their plant’s counterpart during disease development (Fig 3D). A strong positive loading coefficient value (p, 80%) was observed for *RsolAG4-PLR* (Fig 3D) with significant upregulation of transcript abundance in necrotic tissues and surrounding areas by 38.1 fold (*P* = 0.0001) and 34.4 fold (*P* = 0.0001), respectively (Fig 5). *RsolAG4-PDX1* was moderately linked (p, 20%) with necrotic tissue showing an increase of transcript abundance by 1.8 fold (*P* = 0.0004) (Fig 5). However, *RsolAG4-GST* (p, -35%) and *RsolAG4-PDX2* (p, -20%) were negatively associated with necrotic tissue and surrounding areas. (Fig 3C and Fig 3D).

Among the plant-derived genes, *GM-GST* (p, 25%) and *GM-PLR* (p, 15%) were moderately linked with necrotic tissue (Fig 3C and Fig 3D). *GM-PLR* and *GM-GST* showed an increase of 1.5 (*P* = 0.0001) and 1.4 (*P* = 0.0183) folds, respectively, in necrotic tissue (Fig 5). The soybean
GM-CAT encoding gene was positively upregulated in the necrotic tissues with 2.8 ($P = 0.0109$) fold increase (Fig 5). The de novo plant vitamin B6 genes showed an interesting pattern where both GM-PDX2 and GM-PDX1.1 were significantly downregulated by 1.6 ($P = 0.0092$).
Fig 4. Relative transcript abundance of antioxidant genes in pathosystem I. Vitamin B6 (PDX1, PDX2, and PLR), GST, and CAT genes of the pathogen R. solani AG3 (RsolAG3) and the host, Solanum tuberosum (ST) in control and in infected and surrounding tissues at 120 HPI during R. solani AG3 and potato sprouts interaction.
Major plant-derived vitamin B6 genes and GM-GST play a prominent role during soybean RFB disease development caused by R. solani AG1-IA

Upon fungal challenge, the relative transcripts abundance of the soybean PYRIDOXAL REDUCTASE (GM-PLR) and GLUTATHIONE S-TRANSFERASE (GM-GST) was highly upregulated whereas their corresponding fungal genes were notably downregulated (Fig 6). PCA analysis showed that leaf soybean GM-GST (p, 95%), and GM-PLR (p, 20%), were closely linked with the necrotic tissue, with significant increased abundance levels of GM-GST and GM-PLR by 5.3 \( (P = 0.0001) \) and 4.6 \( (P < 0.0001) \) fold, respectively (Fig 6). On the other hand, GM-PDX1.1 (p, -20%) was negatively linked with necrotic tissue and surrounding areas (Fig 3E and Fig 3F), and displayed a significant downregulation with fold changes of 3.4 \( (P < 0.0001) \) and 1.8 \( (P < 0.0001) \) fold, respectively (Fig 6). Although not supported by high loading coefficient values in the PCA loading analysis, the fungal genes RsolAG1-IA-PDX1 (one component of de novo vitamin B6 biosynthesis genes) and RsoAG1-IA-GST exhibited significant down-regulation with fold changes of 2.6 \( (P = 0.0001) \) and 1.5 \( (P = 0.0024) \), respectively, in necrotic tissues, and with fold changes of 3.2 \( (P = 0.0001) \) and 5.6 \( (P < 0.0001) \), respectively, in the surrounding areas (Fig 6).

Accumulation of ROS in R. solani mycelia is related to transcriptional regulation of vitamin B6 machinery and other antioxidant genes

The presence of ROS in the fungal hyphae and plant tissues was detected by fluorescent H$_2$DCF-DA and non-fluorescent DAB methods. Absence of fluorescence was observed in control hyphae grown on half-strength PDA (Fig 7B, Fig 7H and Fig 7N), while an intense green fluorescence was detected in R. solani AG3, AG4, and AG1-IA, hyphae during disease development (Fig 7F, Fig 7L and Fig 7R). Tissues of control and infected potato sprouts, soybean hypocotyls and leaves did not display any green fluorescence (Fig 7D, Fig 7J and Fig 7P) and (Fig 7F, Fig 7L and Fig 7R), respectively.

To demonstrate whether the generation of ROS was promoted by R. solani infection, plant tissues were assayed in situ for the production of H$_2$O$_2$ in response to infection, using DAB staining. DAB oxidizes in the presence of H$_2$O$_2$ generating a dark brown precipitate in plant tissues [41]. Tissues of infected potato sprouts and soybean hypocotyls and leaves displayed a strong brown precipitate specifically at necrotic and surrounding areas (Fig 8B, Fig 8D and Fig 8F). However, control uninfected tissues did not show any browning (Fig 8A, Fig 8C and Fig 8E).

Discussion

It is widely known that ROS management through the use of antioxidants is important for pathogenic microorganisms to prevent excessive oxidative stress [10, 11]. This research further supports our preceding study on the novel function of vitamin B6 genes in R. solani as an antioxidant stress protector against ROS that prevents expansion of oxidative stress [27].
Fig 5. Relative transcript abundance of antioxidants in pathosystem II. Vitamin B6 (PDX1, PDX2, and PLR), GST, and CAT genes of both the pathogen R. solani AG4 (RsolAG4) and the host, Glycine max, (GM) in control and infected and surrounding tissues at 36 HPI during R. solani AG4 and soybean hypocotyls interaction. C: control; R. solani AG4 grown alone or soybean hypocotyl inoculated with sterile PDA. SNL: tissue surrounding necrotic lesion. NL: necrotic lesion.

Asterisks indicate significant relative transcript abundance ratios between the control and the interaction using least significant difference (LSD) test (P < 0.05). Fold change is calculated in relation to the control at 36 HPI. PDX1 and PDX2: Pyridoxine biosynthesis genes, PLR: PYRIDOXAL REDUCTASE, CAT: CATALASE, GST: GLUTATHIONE-TRANSFERASE.

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Necrotrophic fungi can regulate intracellular levels of ROS for developmental and virulence purposes while the host uses ROS to hinder disease progression. A comprehensive understanding of plant responses to fungal pathogens exists [42, 43]. Gene expression studies focusing on oxidative stress responses from both the plant and the pathogen side during plant–R. solani interactions are very limited [28, 34]. This study is the first analysis of gene expression encoding vitamin B6 and several antioxidant genes in plant and fungal tissue of three pathosystems during interaction of different AG groups of R. solani with their respective hosts.

In a previous study, we fully characterized two vitamin B6 de novo biosynthetic pathway genes, RsolPDX1 (KF620111.1) and RsolPDX2 (KF620112.1) genes, and one gene, RsolPLR (KJ395592.1) of the vitamin B6 salvage biosynthetic pathway of R. solani AG3 [27]. The use of mutant yeasts and complementation with PDX1 or PDX2 proved to be a valuable tool in fungi [31, 32]. In this study, we confirmed that R. solani AG3 PDX1 and PDX2 are functionally exchangeable with yeast SNZ1 and SNO1, respectively, and are involved in the de novo vitamin B6 biosynthesis via successful complementation of the S. cerevisiae null mutant Δsnz1 phenotype with RsolAG3-PDX1, and the null mutant Δsno1 with RsolAG3-PDX2. Complementation of each of the S. cerevisiae null mutants (Δsnz1 and Δsno1) with its respective R. solani AG3 gene restored their function implying their ability of de novo production of vitamin B6 in yeast. This also suggests that RsolAG3-PDX1 and RsolAG3-PDX2 can interact with their respective yeast counterpart to form the PDX1–PDX2 complex [44]. Similar to other fungi, our complementation assays validated the role RsolAG3-PDX1 or RsolAG3-PDX2 in oxidative stress resistance through the reversion of the menadione sensitivity of the Δsnz1 or Δsno1 mutant strains, respectively [31, 32]. Taken together, our results demonstrated that R. solani AG3 is probably autotrophic to vitamin B6. However, in order to provide direct evidence of autotrophy to vitamin B6, in vitro assays are required to demonstrate the capability of vitamin B6 synthesis by R. solani AG3 PDX1–PDX2 protein complex. Complementation assays for PDX1/PDX2 of R. solani AG4 and AG1-IA were not attempted, as their respective genes are not characterized.

It is common knowledge that ROS is produced at the site of attempted invasion through an oxidative burst during disease development [9, 10, 28, 45]. Oxidative damage can be prevented in both the host and the pathogen when their antioxidant machinery is efficiently used [12, 46]. Both plants and fungi have ROS scavenging activities that detoxify ROS either directly through the action of enzymes such as: superoxide dismutase (SOD), catalase (CAT), glutathione peroxidase (GPX) and peroxiredoxins (Prx), or indirectly via vitamins like ascorbic acid (vitamin C), tocopherols (vitamin E), and vitamin B6 [23, 46–48].

To describe trends (positive or negative) between the plant’s and fungal antioxidants genes (total 11 antioxidant/pathosystem) in response to stress during plant-pathogen interactions of each pathosystem, it necessitates the use of multivariate analysis. The application of PCA revealed tight groups with no outliers. PCA loading biplots demonstrated that several trends could be observed between the relative transcript accumulation of antioxidant genes and the plant’s areas sampled (necrotic tissues or surrounding areas) in each pathosystem. These results indicate the distinctive differences in regulation of fungal-derived and plant-derived antioxidant genes of each pathosystem in response to R. solani attack of different AGs.
Fig 6. Relative transcript abundance of antioxidants in pathosystem III. Vitamin B6 (PDX1, PDX2, and PLR), GST, and CAT genes of both pathogen *R. solani* AG1-IA (*RsolAG1-IA*) and the host, *Glycine max* (GM) in control and in infected and surrounding tissues at 18 HPI during *R. solani* AG1-IA and...
Our results clearly showed that ROS formation was induced in hyphal cells of *R. solani* AG1-IA, AG3 and AG4 during disease development on their respective hosts. ROS production was also reported in *R. solani* AG 2–2 IV and AG-8, serious pathogens of sugar beet and wheat, respectively [21, 28]. These findings support the notion that several AGs produce ROS. The induced ROS were also linked to the regulation of antioxidant genes and their collective possible role in oxidative stress alleviation. A strong brown precipitate was detected in tissues of infected potato sprouts and soybean hypocotyl and leaves. Accumulation of H$_2$O$_2$ in single cells followed by their death was reported in various plant-fungus interactions [40, 49–51]. These data confirm that *Rhizoctonia* disease development in potato sprouts and soybean hypocotyls and leaves provokes an oxidative stress in fungal hyphae and plant tissues specifically in necrotic areas and could be related to the oxidative status of the fungus as well as the plant.

The differential expression of antioxidant genes in the host and the pathogen is an important determinant of disease outcome and pathogenicity. ROS is under tight control during disease progression where sometimes it can lead to cell death of fungal and plant tissues. Alternatively, pathogen-induced ROS themselves can act as signalling molecule promoting development of fungal structures or resistance in plant cells [11, 28]. In our study, comparison of the differential gene expression across the three pathosystems was not attempted, because the method of inoculation and plant tissue types are different in each pathosystem, resulting in variable infection intensities and different gene expression. Instead, we focused on understanding the expression patterns of fungal and host antioxidant genes in each pathosystem. Our results showed that distinct expression patterns of fungal and host antioxidant genes were associated in necrotic tissues and their surrounding areas for each of the three *R. solani* pathosystems.

During *R. solani* AG3 infection of potato sprouts, significant gene expression of both fungal and plant vitamin B6 *de novo* pathway genes was observed. Genes of the vitamin B6 *de novo*
pathway of *R. solani* and potato (homolog ST-PDX1.1) were highly induced in the surrounding tissue and to a lesser extent in the necrotic tissue. Interestingly, the potato ST-PDX genes (i.e., ST-PDX1.1 and ST-PDX 1.2 homologs of ST-PDX1 and ST-PDX 2) are characterized according to their catalytic ability [52] and were expressed in all potato tissues [37]. The catalytic homolog ST-PDX1.1, but not the non-catalytic homolog ST-PDX 1.2, was induced in response to biotic and abiotic stress (this study, Moccand, Boycheva [52]). These results strongly indicate that ST-PDX1.1 plays a role in disease development. It has been reported that *Arabidopsis* (AT-PDX1.2) acts as a pseudoenzyme that can have a role of a positive regulator of vitamin B6 abundance during abiotic stress [52]. Additionally, it was shown that potato ST-PDX2 might be regulated by ST-PDX1.2, due to the unique (novel) ability of the latter to interact with AT-PDX2 [37, 53]. In our study, the upregulation of ST-PDX2 was not coupled by the upregulation of ST-PDX1.2 during disease development. Whether ST-PDX1.2 acts as a positive regulator of vitamin B6 abundance during potato interaction with *R. solani* merits further studies.

Activation of some of the antioxidant genes was tissue specific. Higher expression of host genes (ST-PDX2, ST-PLR and ST-GST) compared to their counterpart in the pathogen in necrotic tissues was detected. Tissue specificity response is supported by the findings of Denslow, Walls [54]. The increase in the transcript abundance of *PDX1* was evident upon infection of tobacco leaves with *P. syringae pv. phaseolicola* at the region surrounding the infiltration area. However, levels of *PDX1* were lower in the infiltration area where hypersensitive response has developed [54].

In Pathosystem II, both fungal- and plant-derived antioxidant genes played a significant role during disease development where some of them were substantially induced while others were downregulated. Interestingly, fungal and host genes encoding PLR, a member of the
vitamin B6 salvage biosynthetic pathways, were among the driving antioxidant force during soybean hypocotyl- *R. solani* interaction. Both fungal-derived and soybean-derived PLR encoding genes were expressed in the necrotic region as well as in the surrounding tissue, however RsolAG4-PLR was highly expressed. Similarly, in other studies, transcriptional regulation of PLR was substantially induced in response to accumulation of ROS in *R. solani* mycelia parasitized by a mycoparasite or exposed to different abiotic stressors [25, 27, 55]. The results of our study, clearly indicate the involvement of RsolAG4-PLR as an efficient quencher of ROS. On the other hand, the upregulation of soybean PLR (GM-PLR), but not at the same extent as that of the pathogen, in the same areas may imply that the plant may also be limiting ROS damage to the soybean tissue.

The interplay of ROS antioxidant genes produced by fungi and their hosts during their interaction was evident in *R. solani* AG1-IA-soybean leaves interaction (pathosystem III). Major plant-derived antioxidant genes of soybean were upregulated in necrotic leaf tissues whereas the corresponding *R. solani* AG1-IA antioxidant genes showed notable downregulation. These results may possibly indicate that the elevation of antioxidant capacity of two of the three vitamin B6 genes, GM-PDX2 and GM-PLR as well as the enzymatic antioxidant, GLUTATHIONE S-TRANSFERASE (GM-GST) in soybean leaves against *R. solani* AG1-IA increased their tolerance to the development of necrosis and suppressed the spread of the pathogen. These results are supported by the over expression of the antioxidant VB6 genes (i.e., PDX) in *Arabidopsis* that led to increased tolerance to oxidative stress [24, 56]. Equally *Arabidopsis* mutants with defects in vitamin B6 de novo biosynthetic pathway (PDX1.2 or PDX1.3) showed increased levels of disease of gray mold caused by *Botrytis cinerea* [57, 58]. Taken together, these results support the novel function of vitamin B6 genes as antioxidant stress protector against ROS to prevent oxidative stress.

In plant-necrotroph systems, the downregulation of antioxidant genes appears to play an important role during infection and initiation of the oxidative burst, leading to an overflow of ROS in infected areas. For example, in response to infection with *Botrytis cinerea*, leaves of ivy pelargonium (*Pelargonium peltatum*), showed a strong nitric oxide (NO) burst and H$_2$O$_2$ accumulation to arrest disease progression through NO-dependent reversible inhibition of catalase [59]. In our study, the plant’s GmPDX1.1, and ST-CAT in pathosystems I and III were substantially suppressed, respectively, which may lead to enhanced reactive oxygen species generation that typically accompanies infections caused by necrotrophs [60, 61].

**Conclusions**

In conclusion, we confirmed the role of *R. solani* AG3 de novo vitamin B6 genes through yeast complementation assay. In our previous study [27], we provided evidence that vitamin B6 genes of the de novo and the salvage biosynthetic pathways function as antioxidants against oxidative stress induced by ROS chemical inducers [27]. In this study, we extend our research to examine whether genes of vitamin B6 machinery are also implicated as antioxidants in response to oxidative stress formed during plant-pathogen interactions. Our findings present new evidence on: (i) the co-expression of VB6 genes along with other well-known antioxidant genes (i.e., CAT and GST) in both the plant and pathogen side during their interaction, and (ii) the differential transcriptional regulation of the genes in infected necrotic tissues and their surrounding areas of three pathosystems involving different plants and different AGs of *R. solani*.

It is also likely that the requirement of Vitamin B6 genes of the salvage pathway differs among different plant species in response to pathogen infection. It appears that both the pathogen and host employ unique expression strategy depending on the type of interaction as well as the type of tissue affected. Hence, the study offers novel insights into the biological...
correlation and identification of ROS antioxidant genes that can be used in soybean and potato breeding programs for resistance against *R. solani*. Overall, we demonstrate that the co-expression and accumulation of certain plant and pathogen ROS-antioxidant related genes in each pathosystem are highlighted and might be critical during disease development from the plant’s point of view, and in pathogenicity and developing of infection structures from the fungal point of view. Future studies are underway to test whether different potato cultivars containing variable amounts of vitamin B6 [37] exhibit differential degrees of susceptibility to *R. solani* AG 3, and that this susceptibility is correlated with content and differential gene expression of vitamin B6.

**Supporting information**

**S1 Fig.** Principle component analysis score plots (PC1/PC2) for the effect of 11 antioxidant genes relative transcripts abundance on control, necrotic lesions and surrounding areas of necrotic lesions of Pathosystem I (A), II (B) and III (C). The ellipse represents the Hotelling $T^2$ at a 95% confidence interval. Three biological replications were performed per treatment. Q2 (cum); cumulative fraction of the total variation of the X’s that can be predicted by the extracted components, R2X; the fraction of the sum of squares of the two principal components.

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