Spermidine Is Critical for Growth, Development, Environmental Adaptation, and Virulence in *Fusarium graminearum*

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**INTRODUCTION**

Polyamines (PAs) are ubiquitous polycations found in both eukaryotic and prokaryotic organisms. The most common polyamines widely found in all organisms are putrescine (Put), spermidine (Spd), and spermine (Spm) (Igarashi and Kashiwagi, 2019). The Put is produced by ornithine decarboxylase 1 (*SPE1*) from the amino-acid L-ornithine. The synthesis of Spd and Spm is mainly catalyzed by spermidine synthase (*SPE3*) and spermine synthase (*SPE4*), respectively. Both reactions require the previous decarboxylation of S-adenosylmethionine (SAM) which is catalyzed by S-adenosylmethionine decarboxylase (*SPE2*). SAM donates the aminopropyl group for the conversion of Put into Spd, and subsequently of Spd into Spm (Vanrell et al., 2013).

Polyamine biosynthesis (Spe) genes have been cloned and sequenced in *Saccharomyces cerevisiae*. The *SPE1*, *SPE2*, and *SPE3* genes are important for growth in a polyamine-free environment. Polyamine deficiency leads to growth defect even in yeast cells, suggesting that polyamines play a role in growth regulation, development, and stress responses (Zhang et al., 2010; Igarashi and Kashiwagi, 2019). In addition, polyamines are known to influence the expression of genes involved in the metabolism of stress responses (Igarashi and Kashiwagi, 2019).

Putrescine, spermidine, and spermine are the most common natural polyamines. Polyamines are ubiquitous organic cations of low molecular weight and have been well characterized for the cell function and development processes of organisms. However, the physiological functions of polyamines remain largely obscure in plant pathogenic fungi. *Fusarium graminearum* causes Fusarium head blight (FHB) and leads to devastating yield losses and quality reduction by producing various kinds of mycotoxins. Herein, we genetically analyzed the gene function of the polyamine biosynthesis pathway and evaluated the role of the endogenous polyamines in the growth, development, and virulence of *F. graminearum*. Our results found that deletion of spermidine biosynthesis gene *FgSPE3* caused serious growth defects, reduced asexual and sexual reproduction, and increased sensitivity to various stresses. More importantly, Δ*Fgspe3* exhibited significantly decreased mycotoxin deoxynivalenol (DON) production and weak virulence in host plants. Additionally, the growth and virulence defects of Δ*Fgspe3* could be rescued by exogenous application of 5 mM spermidine. Furthermore, RNA-seq displayed that FgSpe3 participated in many essential biological pathways including DNA, RNA, and ribosome synthetic process. To our knowledge, these results indicate that spermidine is essential for growth, development, DON production, and virulence in *Fusarium* species, which provides a potential target to control FHB.

**Keywords:** *Fusarium graminearum*, polyamine biosynthesis, spermidine, DON (deoxynivalenol), virulence
The role of polyamine in the responses to plant stress in *A. thaliana* and *S. cerevisiae* is essential for the survival of these organisms. The gene encoding spermine synthase is also not clearly identified and characterized in *S. cerevisiae* (El-Tarabily et al., 2020, 2021; Mathew et al., 2020). In filamentous fungi, increasing yield, and biomass and reducing abiotic stress effects (El-Tarabily et al., 2020, 2021; Mathew et al., 2020). In filamentous fungi, such as *Aspergillus flavus*, the results showed that inactivation of SPE3 clearly reduced conidiation, aflatoxin production, and virulence on maize seed (Majumdar et al., 2018). Recently, Rocha et al. (2020) illustrated that Spm was essential for development and pathogenicity during the early stages of infection in *Magnaporthe oryzae*. However, the functions of the genes in the polyamine biosynthesis pathway in *Fusarium graminearum* have not been demonstrated. Therefore, further elucidation of the mechanisms underlying polyamine biosynthesis is imperative for the development of novel strategies to manage *F. graminearum*-caused diseases.

Polyamines are required for many fundamental processes of nucleic acid synthesis, the stabilization of RNA, DNA, and protein structure (Igarashi and Kashiwagi, 2000). Put and Spd are predominant polyamines that are often found in fungi, while Spm is absent from some fungal genera (Valdes-Santiago et al., 2012). Polyamine depletion causes a serious growth defect in some organisms, whereas excess polyamines in cells show cytotoxicity, indicating an exact regulation of polyamine metabolism is a necessity for cellular processes (Casero et al., 2018). Additionally, polyamines have been identified to regulate various biological processes, including vegetative growth, spore germination, and stress tolerance and virulence of plant and animal fungal pathogens (Ruiz-Herrera, 1994; Khurana et al., 1996).

_Fusarium head blight* (FHB), also known as head scab disease, is caused by *F. graminearum*, and is a destructive fungal disease of cereal grains worldwide (Dean et al., 2012; Chen et al., 2019). In addition to substantial yield losses, mycotoxin contamination, such as deoxynivalenol (DON), nivalenol, and zearalenone (ZEN), exerts an adverse threat to the health of humans and animals (Chen et al., 2019). DON (also known as vomitoxin) is one of the most commonly occurring trichothecenes in cereals and cereal-derived food products. This can result in feed refusal, immune dysregulation, teratogenicity, and reproductive defects in mammals (Pestka, 2010; Lee and Ryu, 2017). Meanwhile, DON, as a vital virulence factor, plays a vital role in the pathogenic process. It is very important to explore the mechanism of DON synthesis for the control of FHB. DON production was affected by several environmental factors including carbon, nitrogen, pH, and light (Chen et al., 2019). Recently, using a large-scale nutrient profiling, Gardiner et al. (2009a, 2010) revealed that exogenous Put can significantly increase DON production and further testified that Put produced by plants might play a role in inducing DON biosynthesis during the *F. graminearum* infection process. However, the molecular mechanisms in fungi linking the endogenesis polyamine biosynthetic pathway to DON production and pathogenicity are not clear in *F. graminearum*. In this study, four genes (FgSPE1, FgSPE2, FgSPE3, and FgSPE4) which were involved in the polyamine biosynthetic pathway, were identified and genetically analyzed in *F. graminearum*. Further, our study evaluated the function of polyamines, and demonstrated that spermidine was important for hyphal growth, conidial production, DON biosynthesis, environmental stress response, and virulence in this important phytopathogenic fungus.

### MATERIALS AND METHODS

#### Fungal Strains and Growth Assays

The wild-type strain PH-1 (NRRL 31084) was chosen as parental strain for generating all deletion mutants (Cuomo et al., 2007). All strains were cultured on complete medium (CM), potato dextrose agar (PDA), minimal medium (MM), and yeast extract peptone dextrose (YPD) medium at 25°C to assay growth rate.

#### Generation of Gene Deletion and Complementation Strains

The gene deletion mutants were constructed by double-joint PCR method, the primer pairs which were used to amplify the upstream and downstream sequence of the targeted gene are shown in [Supplementary Table 1](#SupplementaryTable1). The resulting products were then transformed into the protoplasts of wild-type PH-1 as described previously (Tang et al., 2020). To complement the ΔFgspe3, the Fgspe3-GFP cassette (GFP, green fluorescent protein) was constructed, a Fgspe3 fragment which included the promoter and gene fragment (without stop codon) was amplified from wild-type genomic DNA with specific primers listed in [Supplementary Table 1](#SupplementaryTable1). The resulting complementation fragments and Xho1 digested pYF11 vector were co-transformed into yeast XK1-25. The recombined Fgspe3-GFP fusion vector was transformed into *Escherichia coli* for amplification and then transformed into the ΔFgspe3 mutant, and the generating

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**Supplementary Table 1**

| Gene | Function |
|------|----------|
| SPE1 | Polyamine biosynthesis |
| SPE2 | Polyamine biosynthesis |
| SPE3 | Polyamine biosynthesis |
| SPE4 | Polyamine biosynthesis |

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**Supplementary Table 2**

| Media | Composition |
|------|-------------|
| CM   | Complete medium |
| PDA  | Potato dextrose agar |
| MM   | Minimal medium |
| YPD  | Yeast extract peptone dextrose |

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**Supplementary Table 3**

| Growth Rate | Description |
|-------------|-------------|
| C            | Control    |
| ΔSPE1       | Deletion    |
| ΔSPE2       | Deletion    |
| ΔSPE3       | Deletion    |
| ΔSPE4       | Deletion    |

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**Supplementary Table 4**

| Deletion Mutants | Description |
|-----------------|-------------|
| PH-1            | Wild-type   |
| ΔSPE1           | Deletion of SPE1 |
| ΔSPE2           | Deletion of SPE2 |
| ΔSPE3           | Deletion of SPE3 |
| ΔSPE4           | Deletion of SPE4 |

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**Supplementary Table 5**

| Complementation Strains | Mutant Description |
|-------------------------|--------------------|
| PH-1                    | Wild-type          |
| ΔFgspe3                 | Deletion of Fgspe3 |
| ΔFgspe3-GFP             | Deletion of Fgspe3-GFP |

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**Supplementary Table 6**

| PCR Primers | Description |
|-------------|-------------|
| Upstream   | Primer for upstream sequence |
| Downstream | Primer for downstream sequence |
| FgSPE1     | Primer for SPE1 gene |
| FgSPE2     | Primer for SPE2 gene |
| FgSPE3     | Primer for SPE3 gene |
| FgSPE4     | Primer for SPE4 gene |

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**Supplementary Table 7**

| Growth Rate | Description |
|-------------|-------------|
| C           | Control    |
| ΔSPE1       | Deletion    |
| ΔSPE2       | Deletion    |
| ΔSPE3       | Deletion    |
| ΔSPE4       | Deletion    |

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**Supplementary Table 8**

| Deletion Mutants | Description |
|-----------------|-------------|
| PH-1            | Wild-type   |
| ΔSPE1           | Deletion of SPE1 |
| ΔSPE2           | Deletion of SPE2 |
| ΔSPE3           | Deletion of SPE3 |
| ΔSPE4           | Deletion of SPE4 |

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**Supplementary Table 9**

| Complementation Strains | Mutant Description |
|-------------------------|--------------------|
| PH-1                    | Wild-type          |
| ΔFgspe3                 | Deletion of Fgspe3 |
| ΔFgspe3-GFP             | Deletion of Fgspe3-GFP |

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**Supplementary Table 10**

| PCR Primers | Description |
|-------------|-------------|
| Upstream   | Primer for upstream sequence |
| Downstream | Primer for downstream sequence |
| FgSPE1     | Primer for SPE1 gene |
| FgSPE2     | Primer for SPE2 gene |
| FgSPE3     | Primer for SPE3 gene |
| FgSPE4     | Primer for SPE4 gene |

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**Supplementary Table 11**

| Growth Rate | Description |
|-------------|-------------|
| C           | Control    |
| ΔSPE1       | Deletion    |
| ΔSPE2       | Deletion    |
| ΔSPE3       | Deletion    |
| ΔSPE4       | Deletion    |

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**Supplementary Table 12**

| Deletion Mutants | Description |
|-----------------|-------------|
| PH-1            | Wild-type   |
| ΔSPE1           | Deletion of SPE1 |
| ΔSPE2           | Deletion of SPE2 |
| ΔSPE3           | Deletion of SPE3 |
| ΔSPE4           | Deletion of SPE4 |

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**Supplementary Table 13**

| Complementation Strains | Mutant Description |
|-------------------------|--------------------|
| PH-1                    | Wild-type          |
| ΔFgspe3                 | Deletion of Fgspe3 |
| ΔFgspe3-GFP             | Deletion of Fgspe3-GFP |

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**Supplementary Table 14**

| PCR Primers | Description |
|-------------|-------------|
| Upstream   | Primer for upstream sequence |
| Downstream | Primer for downstream sequence |
| FgSPE1     | Primer for SPE1 gene |
| FgSPE2     | Primer for SPE2 gene |
| FgSPE3     | Primer for SPE3 gene |
| FgSPE4     | Primer for SPE4 gene |

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**Supplementary Table 15**

| Growth Rate | Description |
|-------------|-------------|
| C           | Control    |
| ΔSPE1       | Deletion    |
| ΔSPE2       | Deletion    |
| ΔSPE3       | Deletion    |
| ΔSPE4       | Deletion    |

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**Supplementary Table 16**

| Deletion Mutants | Description |
|-----------------|-------------|
| PH-1            | Wild-type   |
| ΔSPE1           | Deletion of SPE1 |
| ΔSPE2           | Deletion of SPE2 |
| ΔSPE3           | Deletion of SPE3 |
| ΔSPE4           | Deletion of SPE4 |

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**Supplementary Table 17**

| Complementation Strains | Mutant Description |
|-------------------------|--------------------|
| PH-1                    | Wild-type          |
| ΔFgspe3                 | Deletion of Fgspe3 |
| ΔFgspe3-GFP             | Deletion of Fgspe3-GFP |

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**Supplementary Table 18**

| PCR Primers | Description |
|-------------|-------------|
| Upstream   | Primer for upstream sequence |
| Downstream | Primer for downstream sequence |
| FgSPE1     | Primer for SPE1 gene |
| FgSPE2     | Primer for SPE2 gene |
| FgSPE3     | Primer for SPE3 gene |
| FgSPE4     | Primer for SPE4 gene |
transformant was named as the ΔFgspe3-C. By means of the same methods, the FgsPE1-GFP, FgsPE2-GFP, and FgsPE4-GFP recombinant plasmids were transformed into the mutants, respectively, and generated the ΔFgspe1-C, ΔFgspe2-C, and ΔFgspe4-C complementation strains.

Conidiation, Perithecium Formation Assays
For conidia production, fresh mycelial plugs of each tested strain were inoculated in liquid carboxymethyl cellulose (CMC) medium (0.1% CMC, 2 mM MgSO$_4$, 4 mM NH$_4$NO$_3$, 10 mM KH$_2$PO$_4$, and 1 L water). The 50-mL flasks were cultured in a shaker (180 rpm) for 4 days at 25°C. The number of conidia for each strain in the medium was counted using a hemacytometer (Sigma-Aldrich, United States). The conidial morphology was observed by Calcofluor White (CFW) staining and photographed with a confocal microscope (Gottingen, Niedersachsen, Germany). Perithecium production was inoculated on carrot agar medium (CA) with a 12 h/12 h light/dark cycle at 25°C.

Stress Sensitivity Tests
To determine the growth rate under different stress conditions for deletion mutants, fresh mycelial plugs (5 mm in diameter) of each strain were inoculated on PDA medium containing various stress compounds described in the figure legends. Exogenous NO$_3^−$, NH$_4^+$, Put, Spd, and Spm (Sigma-Aldrich, United States) at 5 mM were separately added into minimum medium minus nitrogen (MM-N) medium to evaluate the growth of the ΔFgspe mutants. MM-N medium which did not include any nitrogen source served as a control.

Pathogenicity and Deoxynivalenol Production Assay
Conidia were collected from a CMC medium and adjusted to 10$^3$ conidia/mL with sterile water. A 10-µL conidial suspension was inoculated into a spikelet of a 6-week-old flowering wheat head. For each strain, 15 replicates were injected into spikelets for analysis. The infected spikelet was characterized 15 days after inoculation. To further analyze the pathogenicity of the ΔFgspe mutants in detail, a fresh mycelia plug of wild-type and mutants was placed on fresh corn silks. Then in 5 days after incubation at 25°C, the extent of discoloration was recorded. For wheat leaf infection assays, the 4-week-old fresh leaves were inoculated with mycelial plugs at 25°C. Pictures were taken 6 days after inoculation. For wheat coleoptile infection assays, 3-day-old seedlings of wheat were used. Conidial suspension was injected and recorded symptoms 7 days after inoculation. For each strain, there were 15 replicates in each experiment.

For detection of DON production, ΔFgspe3 strain was grown in liquid trichoethene biosynthesis induction (TBI) medium at 28°C for 1 week in the dark. The supernatant was harvested by three layers of filter paper and then quantified by a liquid chromatography tandem–mass spectrometry (LC-MS/MS) system (Tang et al., 2018).

Quantitative Real-Time PCR
The wild-type and ΔFgspe3 were cultured into TBI medium for 2 days at 28°C. The mycelia of each strain were harvested, and then total RNA was isolated using the TaKaRa RNAiso Extraction Kit, and then used RevertAid H Minus First Strand cDNA Synthesis Kit to synthesize cDNA (Fermentas Life Sciences, Burlington, ON, Canada). The transcriptional levels of TRJ1, TRJ5, and TRJ6 of PH-1 were determined by quantitative reverse transcription-polymerase chain reaction (RT-qPCR) using the method previously described (Liu et al., 2019). ACTIN (FGSG_07335) gene was served as a control. The primers were listed in Supplementary Table 1.

Confocal Microscopy Observation
The conidia of deletion mutant were stained by 0.1 mg/mL CFW for 30 s and observed with a fluorescent microscope (Gottingen, Niedersachsen, Germany). Hyphal branching patterns were examined with a Nikon microscope.

RNA-Seq Analysis
For deep transcriptome sequencing, 2-day-old mycelia of wild-type and ΔFgspe3 strain were harvested from liquid YEFD shaken at 180 rpm, washed three times, and ground in liquid nitrogen. For each strain, there are three biological replicates independently. Total RNA was isolated, and cDNA libraries were constructed by NEBNext® UltraTM Directional RNA Library Prep Kit (New England Biolabs, Ipswich, MA, United States) using the manufacturer's instructions. The cDNA libraries were sequenced by an Illumina Hiseq Novaseq platform with paired-end reads (150 bp) by Novogene Corporation (Beijing, China). Reads were mapped to the genome of F. graminearum PH-1 strain. DESeq R package was used for differential expression gene analysis by Novogene Corporation (Beijing, China). Genes with log$_2$ Fold change $> 1$ or $<-1$ and $P$-value $< 0.05$ were divided into differentially expressed. Raw data of the RNA-seq experiments generated in this article were deposited in the National Center for Biotechnology Information under the accession number PRJNA759842.

Statistical Analysis
Graphs were plotted by GraphPad Prism 7.0 (GraphPad Software, San Diego, CA, United States). The statistical experiments were evaluated using SPSS Statistics 21.0 software. Statistical analysis was performed one-way ANOVA and Fisher’s LSD test ($P < 0.05$).

RESULTS
Identification of Polyamine Biosynthesis Pathway Genes in Fusarium graminearum
Combined with the other fungi, plants, and animals, we firstly inferred the pathway of polyamine biosynthesis in F. graminearum (Supplementary Figure 1). Further, 1https://www.ncbi.nlm.nih.gov/sra/
using the S. cerevisiae polyamine biosynthesis (Spe) protein sequence to query F. graminearum genome, the heterologous genes involved in polyamine biosynthesis which included FGSG_05903 (designated as Fgspe1), FGSG_09035 (designated as Fgspe2), FGSG_13725 (designated as Fgspe3), and FGSG_09525 (designated as Fgspe4) were revealed in this fungus. To further confirm the Spe protein evolution relationships within the organism, phylogenetic analysis of the Spe protein sequence was performed and indicated that the Spe proteins was highly conserved in fungi. Interestingly, most Spe homologs were notably clustered into one distinct subclade (Figure 1A). Meanwhile, the InterPro database and tools for protein domain analysis indicated that Fgspe protein harbored conserved domains that are unique to Spe proteins (Figure 1B). In general, the domain characteristic and the phylogenetic tree indicated Spe homologs proteins were conserved and might have similar biological functions in other fungi.

**Spermidine Is Important for Hyphal Growth**

Polyamines play an essential role to maintain normal growth and development in a living organism (Igarashi and Kashiwagi, 2019). To further study the function of polyamine biosynthesis gene in F. graminearum, we constructed deletion mutants, ΔFgspe1, ΔFgspe2, ΔFgspe3, and ΔFgspe4, using homologous recombination strategy by replacing open reading frame (ORF) of Fgspe gene with hygromycin B phosphotransferase (hph) gene cassette. Transformants in each mutant showed consistent phenotypes on the same conditions. To verify that the phenotypic defects in mutants were due to gene deletion, we constructed the complemented vector for each mutant and generated the ΔFgspe1-C, ΔFgspe2-C, ΔFgspe3-C, and ΔFgspe4-C complemented strains. As shown in Figures 2A,B, the loss of Fgspe1 resulted in significantly reduced growth in CM medium and MM medium that were polyamine-free mediums. However, the ΔFgspe1 exhibited similar mycelial growth rates compared with the wild-type strain in PDA and YEPD medium that may be included by trace amounts of polyamines. The ΔFgspe2 and ΔFgspe3 mutants showed obviously hyphal growth defects on PDA, CM, MM, and YEPD medium. Moreover, ΔFgspe2 and ΔFgspe3 also showed a reduction in aerial hyphae. However, ΔFgspe4 showed no obvious phenotypic defects. It might indicate that Fgspe4 encoding spermine synthase was not important for the growth.

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**FIGURE 1 | Identification of the Fgspe proteins in F. graminearum. (A) Phylogenetic analysis of SPEs from F. graminearum, S. cerevisiae, and other filamentous fungi. Protein sequences of Fgspe1–4 paralogues were aligned by CLUSTALW, and a neighbor-joining tree was constructed by MEGA 7.0. The names of all proteins are shown in the figure. (B) Schematic drawing of Fgspe superfamily proteins members in F. graminearum. Conserved domains are indicated in the figure.**
FIGURE 2 | Phenotypes of the ΔFgspe mutants in vegetative growth. (A) Colony morphology of wild-type PH-1, the ΔFgspe mutants and complemented strains cultured on PDA, CM, MM, and YEPD medium for 3 days at 25°C. (B) Statistical analysis of hyphal growth rate of wild-type PH-1 and the mutants. The diameter was measured after culturing at 25°C for 3 days. Columns listed with the same letter indicates no obviously difference with the least significant difference (LSD) test at \( P = 0.05 \). (C) The patterns of hyphal branching and tip growth for wild-type and ΔFgspe mutants inoculated in PDB liquid medium for 24 h. Bar, 50 µm. (D) Statistical analysis of biomass of wild-type, the ΔFgspe mutants, and complemented strains for 2 days in YEPD medium.
of *F. graminearum* (Figures 2A,B). Subsequently, microscopic observation exhibited that the hyphal morphology of ΔFgspe2 and ΔFgspe3 mutants were highly branched and misshapen (Figure 2C). The dry weight biomass of ΔFgspe2 and ΔFgspe3 were significantly reduced compared to wild-type in YEPD medium for 2 days (Figure 2D). As shown in Supplementary Figure 1, Fgspe2 and Fgspe3 are essential for Spd synthesis. Therefore, our result indicates that the endogenous Spd is vital for the hyphal growth of *F. graminearum*.

**Polyamines Restore the Defective Phenotypes of ΔFgspe Mutants**

As shown in Figure 3, wild-type PH-1 showed as seriously growth defective in the MM-N medium in which was not included any nitrogen source. To further confirm the function of the polyamine in *F. graminearum* therefore, we firstly testified the growth rate of wild-type PH-1 in MM-N medium by exogenous supplemented polyamine, NO$_3^-$ and NH$_4^+$ served as the control. In Figure 3, the phenotypic defects of wild-type PH-1 in MM-N medium showed better recovery by exogenous supplementation of Spd in comparison with Put and Spm, indicating that Spd exhibits a more important role for the growth of *F. graminearum*. The ΔFgspe1, ΔFgspe2, and ΔFgspe3 mutants showed the seriously defective growth in CM, MM, and MM-N mediums that were free of polyamines (Figures 2A, 3). Put, Spd, and Spm are the downstream metabolites of ornithine decarboxylase (Vanrell et al., 2013). To explore the defective phenotypes of ΔFgspe mutants and whether they could be recovered by the addition of exogenous polyamines, the ΔFgspe mutants were grown on MM-N medium supplemented with Put, Spd, and Spm at 5 mM concentration, respectively. NO$_3^-$ and NH$_4^+$ served as the control. The results showed that NO$_3^-$ and NH$_4^+$ were not conducive for growth restoration of ΔFgspe1, ΔFgspe2, and ΔFgspe3 mutants. When Put was added to MM-N medium, defects in hyphal growth of ΔFgspe1 mutant were restored (Figure 3). When exogenous Spd was added to an MM-N medium, the vegetative growth of the ΔFgspe2 and ΔFgspe3 could be fully rescued. In general, our data further indicates that the Spd plays a vital role in hyphal growth in this fungus.

**Spermidine Is Essential for Asexual and Sexual Reproduction**

Sexual and asexual reproduction play a vital role in the infection cycle of *F. graminearum*. Subsequently, the conidial and perithecia production of ΔFgspe mutants was observed. Firstly, to rule out the effect of growth defects on asexual and sexual reproduction, the growth of ΔFgspe mutants was determined. As shown in Figures 4A,B, the ΔFgspe mutants and complementary strains showed a consistent phenotype in conidiation (CMC) and sexual reproduction (CA) in this medium in comparison with wild-type PH-1. To further confirm the function of the polyamine in *F. graminearum* therefore, we firstly testified the growth rate of wild-type PH-1 in MM-N medium by exogenous supplemented polyamine, NO$_3^-$ and NH$_4^+$ served as the control. In Figure 3, the phenotypic defects of wild-type PH-1 in MM-N medium showed better recovery by exogenous supplementation of Spd in comparison with Put and Spm, indicating that Spd exhibits a more important role for the growth of *F. graminearum*. The ΔFgspe1, ΔFgspe2, and ΔFgspe3 mutants showed the seriously defective growth in CM, MM, and MM-N mediums that were free of polyamines (Figures 2A, 3). Put, Spd, and Spm are the downstream metabolites of ornithine decarboxylase (Vanrell et al., 2013). To explore the defective phenotypes of ΔFgspe mutants and whether they could be recovered by the addition of exogenous polyamines, the ΔFgspe mutants were grown on MM-N medium supplemented with Put, Spd, and Spm at 5 mM concentration, respectively. NO$_3^-$ and NH$_4^+$ served as the control. The results showed that NO$_3^-$ and NH$_4^+$ were not conducive for growth restoration of ΔFgspe1, ΔFgspe2, and ΔFgspe3 mutants. When Put was added to MM-N medium, defects in hyphal growth of ΔFgspe1 mutant were restored (Figure 3). When exogenous Spd was added to an MM-N medium, the vegetative growth of the ΔFgspe2 and ΔFgspe3 could be fully rescued. In general, our data further indicates that the Spd plays a vital role in hyphal growth in this fungus.

![FIGURE 3](image-url) Polyamines restores the growth defection of ΔFgspe mutants. Colony morphology of PH-1, deletion mutants, and complementation mutants in MM-N medium supplemented with different nitrogen sources at 5 mM. Photos were recorded at 25°C after 3 days.
with the PDA medium (Figure 2A). Hence, we further observed the conidial and perithecia production of ΔFgspe mutants in CMC and CA medium. Compared to wild-type, our result indicated that the ΔFgspe2 and ΔFgspe3 exhibited an obvious reduction in conidiation after incubation for 4 days in CMC medium (Figure 4C). To further observe conidial morphology, the conidia of all mutants were stained by CFW dye. The conidial morphology was recorded with a fluorescence microscope. The conidia size of ΔFgspe2 and ΔFgspe3 was shorter in comparison with wild-type (Figure 4D). Moreover, the ΔFgspe2 and ΔFgspe3 mutants could not grow at the high temperature and oxidative stress generated by 30°C and H₂O₂, respectively. ΔFgspe1 shows weakly increased sensitivity to environmental stress in Figure 5A. However, ΔFgspe4 showed a similar susceptibility in the wild-type toward the above stress (Figures 5A, B). The complementation strains exhibited the same sensitivities as wild-type under various stress conditions.

Spermidine Regulates Environmental Stress Tolerance

A previous study conducted by Yamaguchi et al. (2007) has indicated that polyamine is important to the response to various environmental stresses. Therefore, we detected the sensitivities of the ΔFgspe mutants to different environmental stress agents, including sodium dodecyl sulfate (SDS), Congo Red, H₂O₂, and temperature. The sensitivity assays indicated that the ΔFgspe2 and ΔFgspe3 showed significantly increased sensitivity to various stresses compared to the wild-type. Moreover, the ΔFgspe2 and ΔFgspe3 mutants could not grow at the high temperature and oxidative stress generated by 30°C and H₂O₂, respectively. ΔFgspe1 shows weakly increased sensitivity to environmental stress in Figure 5A. However, ΔFgspe4 showed a similar susceptibility in the wild-type toward the above stress (Figures 5A, B). The complementation strains exhibited the same sensitivities as wild-type under various stress conditions. Aurofusarin of Fusarium fungi played an important role in responding to environment adaptation (Xu et al., 2019). Therefore, we observed the aurofusarin production of ΔFgspe mutants and found that ΔFgspe2 and ΔFgspe3 mutants showed increased yellow pigments on PDA medium (Supplementary Figure 2). In general, our results indicate that Spd shows a vital role in the response to abiotic and biotic stresses.
Spermidine Regulates Virulence of *Fusarium graminearum*

To further investigate the function of polyamine in this pathogenic progress, the virulence of the ΔFgspe mutants was evaluated in flowering wheat heads, the leaves of wheat seedlings, corn silk, and wheat coleoptiles. Notably, the wild-type, ΔFgspe1, ΔFgspe4, and complemented strains could develop on the injected spikelets and quickly spread to the neighboring wheat head, causing discoloration symptoms 2 weeks after...
inoculation. However, the ΔFgspe2 and ΔFgspe3 mutants were not capable of penetrating the spikelet (Figure 6A). Additionally, the ΔFgspe2 and ΔFgspe3 demonstrated attenuated virulence on seedling leaves, corn silk, and wheat coleoptiles (Figures 6B–D). To demonstrate that Spd is the main cause of the reduction in virulence of ΔFgspe3, furthermore, we testified whether exogenous Spd supplement assay could rescue the virulence defects of ΔFgspe2 and ΔFgspe3. Our result showed that the addition of exogenous Spd can indeed rescue the virulence defects of the ΔFgspe2 and ΔFgspe3 mutants in planta (Figure 7). These results demonstrate that Spd is important for virulence in F. graminearum.

Spermidine Regulates Deoxynivalenol Biosynthesis

As DON is proposed as a vital virulence factor and assists the fungus spread during F. graminearum infection process (Seong et al., 2009), we guessed that the reduced pathogenicity of ΔFgspe2 and ΔFgspe3 was also partly due to a reduction in DON production. Therefore, the transcriptional levels of TRI genes which are essential for DON synthesis in the ΔFgspe2 and ΔFgspe3 were detected by RT-qPCR assay after incubation for 2 days in liquid TBI medium. These results indicated that the expression levels of TRI6, TRI5, and TRI11 genes were significantly down-regulated in the ΔFgspe2 and ΔFgspe3 compared with wild-type (Figure 8A). Furthermore, the DON synthesis of ΔFgspe2 and ΔFgspe3 mutants showed a dramatic reduction after incubation for 7 days in a TBI medium (Figure 8B). These results suggest that ΔFgspe2 and ΔFgspe3 are essential for DON biosynthesis in F. graminearum.

FIGURE 6 | ΔFgspe2 and ΔFgspe3 mutants have attenuated virulence in plant. (A) Pathogenicity of all tested strains on flowering wheat heads. Wheat heads were photographed at 15 days post-inoculation using a conidial suspension. The injection sites were denoted as black dots. (B) The ΔFgspe2 and ΔFgspe3 mutants also showed significantly reduced virulence on corn silks. Corn silks were inoculated with fresh mycelial plug of each strain and were determined after 5 days post-inoculation at 25°C. (C) Disease symptoms infected by the tested strains on young wheat leaves. The images were photographed at 6 days. (D) Pathogenicity of the wild-type, mutants, and complementation strains to wheat coleoptiles for 7 days.

FIGURE 7 | Exogenous addition of spermidine restores virulence of ΔFgspe2 and ΔFgspe3 mutants in plant. Disease symptom on wheat seedling leaves inoculated with the ΔFgspe2 and ΔFgspe3 mutants which are treated with or without Spd at the final concentration of 5 mM.
**Transcriptional Profiling Analysis of ΔFgspe3**

To further explore the function of Spd, we performed RNA-seq to wild-type and ΔFgspe3 strain in YEPD medium. For each strain, we plotted three biological replicates. For most genes, the replicates produced similar FPKM values, and all points fall near a line with a slope of 1 (Supplementary Figure 3). Transcriptome analysis revealed that 6,957 genes were differentially expressed in the ΔFgspe3 compared with the wild-type, including 3,331 up-regulated and 3,626 down-regulated genes (Figure 9A). Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analyses were performed to identify the function of the differentially expressed genes (DEGs). GO enrichment analysis identified that genes that are significantly expressed in the ΔFgspe3 mutant were mainly involved in biological processes such as organonitrogen compound biosynthetic, peptide metabolic and translation; cellular components such as ribosome, intracellular ribonucleoprotein complex, cytoplasm, and mitochondrion; and molecular functions such as structural constituents of ribosomes, and RNA and ATP binding (Figure 9B). Furthermore, KEGG enrichment analysis showed a list of the top 20 regulated pathways in ΔFgspe3, including ribosome, DNA replication, oxidative phosphorylation, RNA polymerase, and the citrate cycle (TCA cycle) (Figure 9C). Then, we tested transcriptional expression patterns of nine representative DEGs by qRT-PCR, and the trends in gene expression profiles from qRT-PCR were positively correlated with the transcriptomic data, indicating that expression levels based on RNA-seq data were believable (Figure 9D). Taken together, these results indicate that Fgspe3 might modulate the essential cellular process involved in ribosomes, DNA replication, and carbon metabolism in *F. graminearum*.

**DISCUSSION**

As an important class of metabolites, the polyamines are found distributed in almost all organisms and participate in multiple biological processes, including gene expression regulation, translation, stress resistance, autophagy, and pathogenesis (Miller-Fleming et al., 2015; El-Tarabily et al., 2020, 2021; Mathew et al., 2020). In the current study, our data generated from knocking out Fgspe3 agree with previous publications. Majumdar et al. (2018) have reported that Spd is essential for growth, development, the production of aflatoxins, and pathogenesis in *A. flavus*. Consequently, difluoromethylarginine (DFMA) and L-α-difluoromethylornithine (DFMO) inhibitors of ODC and ADC, respectively, can significantly decrease the fungal growth and virulence (Gamarnik et al., 1994). In *F. graminearum*, DFMO could inhibit growth and DON production in vitro. Additionally, some polyamine transport inhibitors were also shown to be effective against *F. graminearum* activity (Crespo-Sempere et al., 2015). In this study, the genes of polyamines biosynthetic pathway have been identified in *F. graminearum*. The data presented in this study are in line with earlier reports on the role of polyamines in *A. flavus* and demonstrate that polyamines are important for growth, development, and stress resistance in *F. graminearum*. Importantly, the metabolic pathways of polyamines are different in plants and fungi. Thus, targeted and precision-regulated polyamines biosynthetic pathway might represent a novel therapeutic strategy to control plant and animal diseases caused by some phytopathogenic fungi (Valdés-Santiago et al., 2012; Amano et al., 2015).

The current study also shows that supplementation of external Spd can extend the lifespan of a range of species, including yeast, flies, and mice. It has been reported that Spd may be an anti-aging vitamin for humans, and deterioration from aging may be associated with a reduction in endogenous Spd.
FIGURE 9 | Transcriptomic analyses between Δfgspe3 and the wild-type PH-1. (A) Volcano plot analysis of differentially expressed genes (DEGs) of Δfgspe3. The x-axis shows the log2fold-change in RNA expression between PH-1 and Δfgspe3, while the y-axis indicates the –log10 of the adjusted P-value for each RNA. The P-values were adjusted using Benjamini–Hochberg multiple testing. The red dots and green dots represent upregulated and downregulated DEGs. (B) Gene Ontology (GO) term enrichment analysis of DEGs of Δfgspe3. (C) Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analysis of DEGs of Δfgspe3. (D) Expression levels of selected genes were detected by RT-qPCR to PH-1 and Δfgspe3 mutant strains in YEPD medium for 48 h. Results are shown relative to PH-1 and are normalized to the housekeeping gene ACTIN. Columns listed with the same letter are not significantly different at P = 0.05.
In many cases, polyamines are most closely associated with stress tolerance (El-Tarabily et al., 2020, 2021; Mathew et al., 2020). The accumulation of endogenous polyamines is crucial for organisms in response to external stress (Shen et al., 2000). Because of the polycationic nature, polyamines may also stabilize the membranes by interacting with membrane phospholipids under stress conditions (Roberts et al., 1986). Polyamines can also enhance stress tolerance that may be due to regulating nutrient balance and production of reactive oxygen species (ROS) through sophisticated signaling systems (Gong et al., 2014). In this study, the ΔFgspe3 mutant was hypersensitive to H2O2 and high temperature. It might indicate that the strong antioxidant capability of polyamines results in the scavenging of ROS on oxidative stress conditions (Minocha et al., 2014). In this study, the ΔFgspe3 mutant showed significantly increased sensitivity to the tested environmental stress compared with wild-type. These results indicate that the polyamine of *F. graminearum*, especially Spd, plays a crucial role in responding to environmental stresses.

In transcription analysis, genes with log2 Fold change > 1 or < −1 and adjusted P-value < 0.05 as the cutoff criterion were selected for subsequent analysis. Our results FgsPE3 participated in many important biological processes such as peptide metabolic, organonitrogen compound biosynthetic, and translation; cellular components such as intracellular ribonucleoprotein complex, intracellular non-membrane-bounded organelle, cytoplasm, and mitochondrion; and molecular function such as structural constituent of ribosome, RNA and ATP binding. In agreement with a previous study (Igarashi and Kashiwagi, 2000), Polyamines are positively charged molecules and thus can interact with intracellular anionic macromolecules, such as DNA, RNA, chromatin, proteins, eukaryotic translation initiation factor 5A (eIF5A), and phospholipids. Thereby they might potentially affect the outcome of several downstream cellular processes, including subsequent changes in the chromatin structure that can affect gene transcription and eIF5A hypusine modification can regulate the protein translation (Watanabe et al., 1991; Rocha and Wilson, 2019). A growing number of studies have indicated that polyamines play a critical role in the methylation and acetylation of histones, RNA, and DNA, and chromatin remodeling (Phadwal et al., 2018; Tamari et al., 2018). Together, these results further testify to our finding that Spd plays vital roles in hyphal growth and development in *F. graminearum*. Further characterization of the functions of those genes of polyamine biosynthetic pathway will help to better understand the roles of polyamines in this fungus.

In the rice blast fungus *M. oryzae*, spermine metabolism was identified to be associated with infection-related morphogenesis via redox buffering of the ER underpinning appressorial adhesion and rice cell invasion (Rocha et al., 2020). Another role of polyamines was also characterized in FHB caused by *F. graminearum* (Gardiner et al., 2010). Interestingly, the production of DON is significantly induced by several plant metabolites including Put, ornithine, and arginine which are the main natural precursors of polyamines synthesis (Gardiner et al., 2009a). However, the Fgspe4, encoding spermine synthase, showed no obvious defective phenotypes including growth, development, and pathogenesis. These results show that the functions of polyamine vary dramatically among fungal species. We found, in this study, that ΔFgspe2 and ΔFgspe3 mutants were non-pathogenic. The DON production level of ΔFgspe2 and ΔFgspe3 mutants were significantly reduced in comparison with wild-type. DON is indispensable for the pathogen to extend from infected head to healthy head (Gardiner et al., 2009b).

It may be indicated the endogenous spermidine is critical for infection of *F. graminearum*. Additionally, the slower growth of the ΔFgspe2 and ΔFgspe3 mutants may be partially responsible for the reduced virulence in plants. Whether the reduction in pathogenicity in ΔFgspe2 and ΔFgspe3 mutants during infection is due to the decrease in Spd or possible alteration in relative DON content or both, will require further investigation. Overall the results suggest polyamine biosynthesis process is an attractive target to control FHB and reduce DON contamination.

**DATA AVAILABILITY STATEMENT**

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: https://www.ncbi.nlm.nih.gov/, PRJNA759842.

**AUTHOR CONTRIBUTIONS**

GT, ZM, and WL: conceptualization, data analysis, and writing the manuscript. GT, HX, and JL: experiments. All authors have read and agreed to the published version of the manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2021.765398/full#supplementary-material

**Supplementary Figure 1** | Polyamines biosynthetic pathway.

**Supplementary Figure 2** | Comparisons of pigment biosynthesis of wild-type and ΔFgspe mutants generated in this study. Photos were recorded after each strain was grown for 7 days on a PDA medium.

**Supplementary Figure 3** | Global transcriptional analysis. (A) Biological replicates are highly correlated. Boxplot represents FPKM distributions of different samples. The y-axis is log2 (FPKM + 1). (B) Gene expression levels are displayed at different FPKM levels. The x-axis indicates log2 (FPKM + 1), and the y axis shows the density of the gene.
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