TaUGT6, a Novel UDP-Glycosyltransferase Gene Enhances the Resistance to FHB and DON Accumulation in Wheat

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Fusarium head blight (FHB), a devastating wheat disease, results in loss of yield and production of mycotoxins including deoxynivalenol (DON) in infected grains. DON is harmful to human and animal health and facilitates the spread of FHB symptoms. Its conversion into DON-3-glucoside (D3G) by UDP-glycosyltransferases (UGTs) is correlated with FHB resistance, and only few gene members in wheat have been investigated. Here, Fusarium graminearum and DON-induced TaUGT6 expression in the resistant cultivar Sumai 3 was cloned and characterized. TaUGT6::GFP was subcellularly located throughout cells. Purified TaUGT6 protein could convert DON into D3G to some extent in vitro. Transformation of TaUGT6 into Arabidopsis increased root tolerance when grown on agar plates containing DON. Furthermore, TaUGT6 overexpression in wheat showed improved resistance to Fusarium spread after F. graminearum inoculation. Overall, this study provides useful insight into a novel UGT gene for FHB resistance in wheat.

Keywords: wheat, UDP-glucosyltransferase, resistance, Fusarium head blight, deoxynivalenol

INTRODUCTION

Wheat (Triticum aestivum L.) is the third largest grain crop after corn and rice in terms of production, with > 700 million tons consumed by 2.5 billion people worldwide (Ma et al., 2020). However, wheat production is challenging due to the increasing threat of abiotic and biotic stresses. Fusarium head blight (FHB), primarily caused by Fusarium graminearum, is one of the most devastating wheat diseases in China and other parts of the world, where frequent rainfall occurs from the flowering to early grain filling stages (Ma et al., 2019). This disease can cause marked yield loss via poor grain filling and reduced test weight, with up to 70% yield losses under favorable conditions (Bai et al., 2000). More importantly, FHB-infected grains may contain high concentrations of mycotoxins such as deoxynivalenol (DON) produced by Fusarium species, which are harmful to human beings and animals and have become an even more critical public concern (Ma et al., 2019). Furthermore, DON is considered a virulence factor that induces FHB spread within a spike, and DON delivery to the rachis node could be a trigger for biotrophy–necrotrophy switch (Lemmens et al., 2005; Bönnighausen et al., 2019).
**Fusarium** head blight resistance is a complex quantitative trait controlled by multiple quantitative trait loci (QTLs), and approximately 500 QTLs have been identified since 1999 when the first FHB resistance QTL was published for wheat (Bai et al., 1999; Waldron et al., 1999; Buerstmayr et al., 2019). Five FHB-resistant QTL types were reported in wheat, e.g., type I representing resistance to initial infection, type II representing resistance to fungal spread, and type III representing resistance to mycotoxin accumulation (Schroeder and Christensen, 1963; Mesterházy, 1995). *Fhbl*, a major QTL for type II resistance, is the most comprehensively studied region to date and was cloned by different groups (Rawat et al., 2016; Li et al., 2019; Su et al., 2019). However, QTLs related to other resistance types have not been fully elucidated, and only few type III resistance QTLs have been tested for their ability to detoxify DON or enhance resistance to this toxin (Gunupuru et al., 2017). Cloning genes related to DON resistance or detoxification of transgenic or gene-edited crops may provide an important method to improve FHB resistance in wheat.

Deoxynivalenol is an aggressiveness factor with an important role in spike and seed colonization, and its detoxification will help reduce pathogen virulence and increase FHB resistance, leading to fewer symptoms related to DON-susceptible wheat cultivars (Lemmens et al., 2005). Many genes reportedly affect DON resistance in a direct manner, including genes encoding transporters, detoxification enzymes, and regulators of hormone signaling (Gunupuru et al., 2017). The ABC transporters *PDR5* and *TaABC3.1* were reported to contribute to DON tolerance (Mitterbauer and Adam, 2002; Walter et al., 2015). The maintenance of hormone homeostasis may contribute to DON tolerance. Silencing of the *ethylene insensitive 2* gene in wheat resulted in FHB resistance and gain of function related to gibberellin acid-sensitive DELLA NIL lines that showed more resistance to the *Fusarium*-infected spikes and reduced DON-induced cell death rate (Chen et al., 2009; Saville et al., 2012). Enzymes involved in diverse processes have been shown to enhance DON resistance, e.g., the wheat cytochrome P450 *TaCYP72A*, which contributes to DON host resistance and bacterial cytochrome P450 *Ddha*, which can convert DON to 16-hydroxy-DON (Ito et al., 2013; Gunupuru et al., 2018). UDP-glycosyltransferases (UGTs) are the most widely studied enzymes that have been shown to confer DON and FHB resistance via conjugation of DON to DON-3-O-glucoside (D3G).

UDP-glycosyltransferases are a multigenic and highly divergent superfamily of enzymes widely found in all living organisms, many of which share a close relationship with disease resistance (Gauchon et al., 2005). The C-terminus of UGTs contains a conserved region of 44 amino acids, which is termed plant secondary product glycosyltransferase (PSPG) (Vogt and Jones, 2000). Many UGTs were found based on this conserved region such as >100 UGTs found in wheat (He et al., 2018). The first identified UGT capable of detoxifying DON was *DOG1*, which catalyzed the transfer of glucose to DON, creating a less toxic D3G, and overexpressing this gene results in increased DON tolerance in *Arabidopsis* (Poppenberger et al., 2003). Moreover, other UGTs have been reported to be involved in DON tolerance or FHB resistance. Barley *HvUGT13248* is highly induced by DON and can convert DON to D3G, and the overexpression of this gene in *Arabidopsis* and wheat reduced sensitivity to DON and FHB spread (Schweiger et al., 2010; Li et al., 2015). Two homologs of *HvUGT13248* were found in *Brachypodium distachyon*, and overexpression of *Bradi5g03300* increased root tolerance to DON and enhanced FHB resistance (Schweiger et al., 2013a; Pasquet et al., 2016). *HvUGT10W1* from an FHB-resistant barley cultivar also contributes to FHB resistance (Xing et al., 2017). In wheat, only few UGT genes have been identified. *TaUGT1*, *TaUGT2*, and *TaUGT4* were only analyzed at the transcriptional level following *F. graminearum* infection and were not further characterized (Lin et al., 2008; Ma et al., 2015). *TaUGT12887/TraesCSSB02G148300* conferred increased DON resistance when expressed in yeast (Schweiger et al., 2013b). *TaUGT3* and *TaUGT5* can enhance DON tolerance when expressed in both *Arabidopsis* and wheat (Ma et al., 2010; Xing et al., 2018; Zhao et al., 2018). The expression of *Traces_2BS_14CA35D5D*, orthologous to *Bradi5g03300*, is highly inducible by DON, and transformation of this UGT-encoding gene from wheat into *B. distachyon* enhanced FHB resistance and root tolerance to DON (Gatti et al., 2018). In this study, we cloned and characterized a novel *UGT* gene *TaUGT6* involved in resistance to DON tolerance and *Fusarium* spread.

### MATERIALS AND METHODS

#### Plant Materials and Growth Conditions

Sumai 3, Annong 8455, and Fielder were grown in plastic pots or in single-row-plots in greenhouses at campus of Jiangsu Academy of Agricultural Sciences, Nanjing, China. Spikelets at the early anthesis stage were chosen for *F. graminearum* or DON inoculation, as described previously (Xu et al., 2004). Ten microliters of *F. graminearum* conidia suspension (approximately 5 × 10⁵ conidia spores/mL) were injected into one floret at intermediate positions of spikelets and covered with plastic bags for 3 days to retain moisture. The infected spikelets were harvested at 12, 24, 48, and 72 h after inoculation. During DON inoculation, 5 μL DON (10 mg/L) (Sangon, China) was injected into florets and collected after 2, 4, 8, 12, and 24 h of inoculation. All samples were frozen immediately in liquid nitrogen and stored at -80°C. *A. thaliana* were grown in a chamber under a 16/8-h light/dark condition at 21°C.

#### RNA Extraction and RT-qPCR

Total RNA was isolated using the Promega SV total RNA isolation system (Promega, United States), and first-strand cDNAs were synthesized using the PrimeScript 1st strand cDNA Synthesis Kit (Takara Bio, Dalian, China). RT-qPCR was performed using a Roche thermal cycler 96 using the SYBR Green reagent (Takara Bio, Dalian, China). *Tubulin* in wheat and *actin* in *A. thaliana* were used as internal reference genes. RT-qPCR was performed as described previously (He et al., 2018). Triplicates were applied for all qPCR analyses in this study. Data were calculated using the 2^-ΔΔCt_ method (Livak and Schmittgen, 2000). The primers used in RT-qPCR are listed in Supplementary Table 1.
Subcellular Localization of TaUGT6

Full-length cDNA of TaUGT6 was cloned at the Xho I and Spe I sites of the pA7-eGFP plasmid with the primers listed in Supplementary Table 1. The resultant plasmids were coupled with gold particles, bombarded into onion epidermal cells, and observed as previously described (Scott, 1999). Fluorescence signals of green fluorescent protein (GFP) were observed under UltraView VOX (PE, United States).

Sequence Alignment and Phylogenetic Analysis

TaUGT6 and several other UGT proteins (TaUGT1, TaUGT2, TaUGT3, TaUGT4, TaUGT5, HvUGT13248, Bradi5g03300, and DOGT1) that had been previously reported to contribute to FHB resistance were selected for alignment by DNAMAN using the pairwise method (Poppenberger et al., 2003; Lin et al., 2008; Ma et al., 2015; Pasquet et al., 2016; Li et al., 2017; Xing et al., 2018; Zhao et al., 2018). BLAST analysis of the full-length amino acid sequences of TaUGT6 was performed to identify close homologs of TaUGT6 in the Phytozome database. Multiple sequence alignment of UGT protein sequences was performed by MUSCLE and adjusted manually using GeneDoc. Neighbor-joining trees based on the full-length UGT protein sequences were constructed using MEGA 7.0 with the following parameters: Poisson correction, pairwise deletion, and bootstrap (1000 replicates; random seed).

Glucosyltransferase Activity Assay

Full-length cDNA of TaUGT6 was cloned into a prokaryote expression plasmid pGEX-4T-1, and the GST-tagged TaUGT6 recombinant protein was expressed in E. coli and purified using affinity chromatography (Liu et al., 2020). Then, glucosyltransferase activity assay was performed using a reaction mix (200 μL) containing 20 μL of 0.5 M Tris-HCl (pH = 7.0), 10 μL of 50 mM MgSO4, 10 μL of 200 mM KCl, 5 μL of 0.1 M UDP-glucose, 2 μL of 10% (v/v) β-mercaptoethanol, 5 μL of 10 mM DTT, and 2 μg of the recombinant protein. Following incubation for 3 h at 30°C, 20 μL of 240 mg/ml trichloroacetic acid solution was added to the reaction mix. Then, the reaction mix was filtered with a 0.22-μm membrane, frozen immediately in liquid nitrogen, and stored at −80°C until used for high-pressure liquid chromatography/electrospray ionization–tandem mass spectrometry (HPLC-MS/MS).

Generation of Transgenic Wheat and FHB Resistance Evaluation

TaUGT6 was cloned into the vector pTCK303 with maize ubiquitin promoter. pTCK303–TaUGT6 was transformed into the A. tumefaciens strain EHA105 and then into Fielder, which was highly susceptible to FHB by the Agrobacterium-mediated method (Zhang et al., 2018). Transgenic plants were selected through hygromycin resistance evaluation and PCR. The independent T3 homozygous lines TaUGT6-2, TaUGT6-6, and Col-0 seeds were sterilized and placed on 1/2 MS solid medium for a 2-day dark treatment at 4°C to synchronize germination and then for 3 days in a chamber under a 16/8-h light/dark cycle at 21°C. The seedlings with the same size and growth vigor were transferred to 1/2 MS media containing different concentrations of DON (5 and 30 ppm). Seedlings under 5 and 30 ppm DON treatments were grown for 7 and 21 days, respectively, under identical environmental conditions before the phenotype was documented. The longest root of each seedling was recorded, and the significant level of data was analyzed using Student’s t-test.

Mycotoxin Extraction

Freshly wheat spikes were grounded to powder at 15 dai, shaken with acetonitrile/water (1:1, v/v), which had been treated with 0.1% methanoic acid (ROE, United States). Then, it was centrifuged for 5 min at 3,500 g, and 2 ml of the supernatant was purified with an extraction salt pack (200 mg of magnesium sulfate, 100 mg of sodium citrate, 100 mg C18, and 100 mg of primary secondary amine). After centrifugation, the supernatant was transferred to a new tube and concentrated under nitrogen. Before analysis, the extracts were diluted with 0.6 mL extract solution, filtered using 0.22 μm filters, and then were analyzed by HPLC-MS/MS.

HPLC-MS/MS

The procedure of the HPLC-MS/MS was performed as described by Qiu et al. (2020). Briefly, these analyses were analyzed with a LC-20ADXR liquid chromatograph (Shimadzu, Japan) coupled to an AB SCIEX TRIPLE QUAD 3500 triple-quadrupole mass-spectrometric detector (Applied Biosystems, United States). A Kinetex 100A C18 column (100 × 4.6 mm, 2.6 μm) from Phenomenex (Torrance, United States) was used. Of each sample, 2 μL was injected into the column, the flow rate was 0.5 mL/min, and the column temperature was maintained at 40°C. The mobile phase consisted of 5 mM ammonium acetate-acetic acid (99.9/0.1, v/v) (A) and methanol (B). Gradient elution was performed with the following conditions: 0–0.01 min with a
Scanning Electron Microscopy
The infected spikelets at 3 days after inoculation (dai) were fixed in 3% glutardialdehyde for 2 h, dehydrated in a graded alcohol series, coated with gold and examined using Zeiss EVO LS10 scanning electron microscope (Carl Zeiss, Germany).

RESULTS
Inducible Expression of TaUGT6 by *F. graminearum* and DON in the FHB-Resistant Wheat Cultivar Sumai 3
In a previous study, we conducted genome-wide analysis of family-1 UDP-glycosyltransferases in wheat and found many UGT genes that were upregulated after *F. graminearum* inoculation (He et al., 2018). The novel gene, TRIAE_CS42_5BL_TGACv1_404184_AA1288920, termed TaUGT6, was chosen for further characterization in the present study. We firstly performed quantitative reverse transcription polymerase chain reaction (RT-qPCR) to determine the expression level of TaUGT6 following *F. graminearum* inoculation of spikes at different time points in the FHB-resistant cultivar Sumai 3 and susceptible cultivar Annong 8455. TaUGT6 transcript level was induced at 48 h after *F. graminearum* inoculation (hai) and rapidly increased at 72 hai in Sumai 3. This increase was much lower in Annong 8455 (Figure 1A). TaUGT6 expression level following DON treatment was further analyzed in both Sumai 3 and Annong 8455. TaUGT6 transcript level gradually increased 2 h after DON inoculation (hai) after reaching a peak at 8 hai in Sumai 3, whereas was hardly induced in Annong 8455 treated with DON (Figure 1B). Several UGT proteins that had been previously reported to contribute to FHB resistance were aligned by DNAMAN with the pairwise method. The results revealed that TaUGT6 shared a low sequence similarity with TaUGT1/3 (32.94%), TaUGT2 (35.08%), TaUGT4 (22.36%), and TaUGT5 (25.54%) at the amino acid level, respectively. However, all comprised a 44-amino acid UGT consensus sequence, also referred to as PSPG.

Sequence Analysis of TaUGT6
Based on the public Chinese Spring sequence, full-length TaUGT6 cDNA encoding a predicted protein of 490 amino acids was amplified. The 1200-bp DNA sequence upstream of TaUGT6 and the open reading frame were amplified in both Sumai 3 and Annong 8455, and no differences were observed between the two cultivars. The protein sequences of several UGT proteins that had been previously reported to contribute to FHB resistance were aligned by DNAMAN with the pairwise method. The results revealed that TaUGT6 shared a low sequence similarity with TaUGT1/3 (32.94%), TaUGT2 (35.08%), TaUGT4 (22.36%), and TaUGT5 (25.54%) at the amino acid level, respectively. However, all comprised a 44-amino acid UGT consensus sequence, also referred to as PSPG.
Subcellular Localization of TaUGT6

The plasmid pA7-TaUGT6-eGFP was delivered into onion epidermal cells via gene gun to determine the subcellular localization of TaUGT6; pA7-eGFP was used as a control. Fluorescence observation revealed that TaUGT6 was distributed throughout cells including in cell membrane and nuclei, similar to that in pA7-eGFP (Figure 3). Its subcellular localization was similar to that reported for TaUGT3 but different from signature box at the C-terminus (Supplementary Figure 1) (Vogt and Jones, 2000). A neighbor-joining phylogenetic tree was constructed using MEGA7.0, and the results showed that apart from Arabidopsis DOGT1, TaUGT6 showed high relation to that of known glycosyltransferases, which indicated that they might have a conserved function (Figure 2A).

To further analyze the phylogenetic relationship between TaUGT6 and its close homologs, basic local alignment search tool (BLAST) searches were conducted using the full-length amino acid sequence of TaUGT6 in the Phytozone database, and an unrooted tree was constructed using the neighbor-joining method (Figure 2B). Close homologs of TaUGT6 were found in some grass species, with the closest homolog being barley HORVU5Hr1G104580.1 (Figure 2B).

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TaUGT6 Catalyzes DON Glucosylation in vitro

TaUGT6 encodes a putative UGT, and its activity has not been previously demonstrated. We then tested its glucosyltransferase activity in vitro. Enzyme activity analysis was conducted using UDP-glucose as the sugar donor and DON as the substrate, together with purified GST or GST-tagged TaUGT6 proteins, followed by HPLC-MS/MS. The standards DON and D3G were clearly found at 3.15 min and 3.10 min with m/z 297.200/203.100 for DON (Figure 4A), and m/z 503.100/427.100 for D3G (Figure 4B). D3G was not detected in the reaction mixture containing GST (Figure 4D), but could be observed in the reaction mixture containing GST-tagged TaUGT6 after 3 hours reaction (Figure 4F), which indicated that TaUGT6 was able to transform DON into the less toxic form D3G to some extent (Figure 4E). D3G exhibited a dramatically reduced ability to inhibit protein synthesis and other damage during Fusarium infection, which suggested a potential role of TaUGT6 in FHB resistance.

TaUGT6 Overexpression Increases Root Tolerance to DON

DON reportedly affected root growth on an agar medium (Ma et al., 2010; Pasquet et al., 2016). To determine the involvement of TaUGT6 and the effects of DON on root growth, we firstly made a construct that allowed the constitutive expression of TaUGT6 under the control of the tandem 35S promoter, which was transformed into Arabidopsis thaliana by the floral dip method. T3 homozygous seeds that had a high expression level of TaUGT6 from two independent overexpressing transgenic lines (Figure 5A), i.e., TaUGT6-2 and TaUGT6-6, and wild-type (WT) Col were germinated on toxin free medium, and 3-days-old seedlings with the same size and growth vigor were transferred to agar containing 5 or 30 ppm DON. Following 7 days of growth, the phenotype of Arabidopsis treated with 5 ppm DON was analyzed. The main toxic effects observed were root growth inhibition (Figure 5B). The transgenic lines showed low inhibitory levels, with the average values for the longest root length in TaUGT6-2 and TaUGT6-6 being 1.47 and 1.5 cm, respectively, compared with 1.13 cm in Col (Figures 5B,C). Root growth was severely inhibited after 21 days of 30 ppm DON exposure, and the longest root lengths of TaUGT6-2 and TaUGT6-6 were significantly greater than that of Col.
TaUGT6 Overexpression Enhances Resistance to Fusarium Spread

Investigation of the impact of TaUGT6 on *F. graminearum* infection involved the following. We first transformed *TaUGT6* with a maize ubiquitin promoter into susceptible cultivar Fielder via the *Agrobacterium*-mediated method. Two independent overexpressing transgenic lines TaUGT6-779 and TaUGT6-790 with higher basal transcript level (untreated) than the WT line were chosen for further analysis (Figure 6A). The resistance level of the T3 lines TaUGT6-779 and TaUGT6-790 and the WT Fielder was tested by a single floret inoculation and recorded 7 days after inoculation (dai). No significant difference was found in terms of the proportion of symptomatic spikelet (PSS) between Fielder and the transgenic lines at 7 dai, but TaUGT6 overexpression lines clearly displayed the phenotype less (Figure 6B and Supplementary Figure 2). Furthermore, the overexpressing transgenic lines grown in single-row-plots in the greenhouse showed significantly low PSS from 9 to 21 dai (e.g., the average PSS in TaUGT6-779 and TaUGT6-790 was 36.5% ± 11.9% and 39.1% ± 12%, compared with 55.7 ± 22.3% in Fielder at 21 dai, respectively) (Figures 6B, C). Similar results were found in the overexpression lines grown in plastic pots (Figures 5D, E), which indicated that *TaUGT6* overexpression could increase root tolerance to DON accumulation.
FIGURE 7 | Scanning electron micrographs of wheat spike colonization by F. graminearum. (A) Hyphal network on the inner surface of the lemma in wild-type Fielder (WT) 3 days after inoculation (dai), with a partial enlargement of the right side (red box). (B) Hyphal network in the TaUGT6-779 at 3 dai. (C) Hyphal network in the TaUGT6-790 at 3dai. Bars = 100 \mu m in the left, and 10 \mu m in the right.

in the greenhouse (e.g., the average PSS in TaUGT6-779 and TaUGT6-790 was 48.5 ± 14.3% and 46.4 ± 25.4%, compared with 66.1 ± 11.6% in Fielder at 15 dai, respectively) (Supplementary Figure 2). For further exploration of the FHB resistance in the transgenic lines, the infection condition of F. graminearum in spikes was investigated using scanning electron microscopy. Three days after inoculation, the fungus had developed a dense hyphal network on the inner surfaces of the lemma in the WT Fielder (Figure 7A). However, a less dense hyphal network was clearly seen in both TaUGT6-779 and TaUGT6-790, and the hyphal was not smooth in the WT Fielder (Figures 7B,C). These results showed that TaUGT6 overexpression could increase FHB resistance in plant tissue.

TaUGT6 Overexpression Decreases the Contents of DON

To test whether the overexpression of TaUGT6 reduced DON accumulation, DON and D3G contents were measured 15 dai in whole infected spikes in WT and transgenic lines grown in single-row-plots in the greenhouse by HPLC-MS/MS. The DON contents of the lines TaUGT6-779 and TaUGT6-790 were significantly lower than that of WT (P < 0.01) (Figure 8A). The quantity of total DON (DON+D3G) was reduced to 41% and 61% in lines TaUGT6-779 and TaUGT6-790, respectively, in comparison with the control (Figure 8A). Furthermore, the relative abundance of D3G in “total DON” - D3G/(DON+D3G) was 5.18% and 5.26% for TaUGT6-779 and TaUGT6-790, and 3.98% for WT on a molar basis, respectively, indicating a slight but significant increase in the transgenic lines (Figure 8B).

DISCUSSION

Breeding to improve FHB resistance, particularly DON accumulation resistance, was the most reliable strategy to control FHB and minimize both yield and quality losses, and knowledge regarding the genetic basis underlining wheat–F. graminearum interaction was incorporated in this study. In the past few years, massive efforts have been made to fight against FHB such as successful cloning of Fhb1 genes and application of this QTL into practical breeding as well as production of improved cultivars (Buerstmayr et al., 2019). However, the genetic basis of other resistance types, particularly detoxification of DON or enhanced resistance to the toxin, has not yet been elucidated (Gunupuru et al., 2017). Recently, DON glucosylation, catalyzed by UGTs, has drawn the attention of researchers as a potential source of FHB resistance in plants. Arabidopsis DOGT1 was the first identified UGT capable of detoxifying DON, and barley HvUGT13248 and its homolog Bradi5g03300 in B. distachyon were shown to reduce sensitivity to DON and limit FHB spread (Poppenberger et al., 2003; Li et al., 2015; Pasquet et al., 2016). In wheat, few UGT genes have been identified and proven to be functional in FHB resistance. TaUGT1 was repressed, and TaUGT2 and TaUGT4 with high similarity to HvUGT13248 were induced after F. graminearum inoculation,
but no further functional analysis was conducted (Lin et al., 2008; Ma et al., 2015). Thus far, only TaUGT3, TaUGT5, and Tra02_2BS_LCAASD5D5 (orthologous to Bradi5g03200) in wheat have been transformed into plants to prove their role in FHB resistance and DON tolerance (Gatti et al., 2018; Xing et al., 2018; Zhao et al., 2018). Thus, discovering more UGT genes involved in FHB resistance will help elucidate the host resistance mechanisms that confer resistance to DON and enhance DON detoxification mechanisms and will promote the development of more strategies for improving FHB resistance.

Based on our previous phylogenetic analysis, TaUGT6 was found to be a family 1 UGT Group P member, whereas the previously identified members of TaUGT1, TaUGT2, and TaUGT3 belonged to Group D and TaUGT4 and TaUGT5 belonged to Groups L and E, respectively (He et al., 2018; Zhao et al., 2018). Close homologs of TaUGT6 were only found in grass species, and the closest homolog was barley HORVU5Hr1G104580.1. Moreover, except the conserved PSPG motif at the C-terminus, TaUGT6 exhibited a low sequence similarity with TaUGT1/3 (32.94%), TaUGT2 (35.08%), TaUGT4 (22.36%), and TaUGT5 (25.54%) at the amino acid level, respectively. TaUGT6 is a novel UGT member that might be involved in FHB resistance. TaUGT6 was highly induced after F. graminearum or DON treatment in Sumai 3, whereas this was not the case in Anmogn 8455. Furthermore, no genetic variation was found within the promoter and gene body regions of TaUGT6 between Sumai 3 and Anmogn 8455. The different expression pattern after F. graminearum or DON treatment in these two cultivars might be caused by epigenetic differences such as high DNA methylation in Anmogn 8455, which is the subject of our ongoing research. Many studies have shown that epigenetic mechanisms have been involved in fine tuning the responses of wheat to environmental stresses (Kong et al., 2020). For example, due to epigenetic events, TaCYTP81D5 conferred salinity tolerance in salinity-tolerant cultivars; however, this was not the case with sensitive cultivars despite their similar sequences (Wang et al., 2019). Plant UGTs have different subcellular locations such as TaUGT3 localized in the plasma membrane and nuclei (Ma et al., 2010), TaUGT4 in the cytoplasm (Ma et al., 2015), and TaUGT5 in the plasma membrane (Zhao et al., 2018). TaUGT6 localized throughout the cells including plasma membrane and nuclei. The existence of TaUGT6 in the cellular membrane is supposed to play a role in prohibition of DON binding to the cellular membrane in the DON resistant wheat varieties or enhancing the tolerance to DON by retarding fungal colonization (Snijders and Krecting, 1992; Ma et al., 2010; Zhao et al., 2018). The function of TaUGT6 localized in the nucleus is still unknown, but further research is needed to clarify whether increased detoxification leads to more residual ribosomal activity allowing better expression of induced defense transcripts into proteins executing FHB resistance or TaUGT6 influence/interact with other proteins in the nuclei to execute the function of FHB resistance and other processes.

The availability of genomic data has been made more convenient to identify new plant UGTs. However, it was difficult to determine which UGTs possessed DON-inactivating capabilities according to phylogenetic-based homology (Osmani et al., 2009). For example, many homologous UGTs of AtUGT73C5 or HvUGT13248 from rice, barley, and B. distachyon showed no capability of DON detoxification (Schweiger et al., 2013a). Earlier determination of the catalytic activity of UGTs by heterologous expression in E. coli or yeast and in vitro enzyme activity analysis saved time during the identification of new functional UGTs. In this study, we analyzed enzymatic activity with purified GST-tagged TaUGT6 proteins before conducting genetic work. The catalytic activity (0.25 mM DON/2.5 mM UDP-glucose) of TaUGT6 was carried out 3 hours after reaction, and D3G was detected. Determining the optimum reaction conditions may get a better catalytic efficiency. TaUGT6 was able to transform DON into the less toxic form D3G to some extent in vitro, which suggested that it might play a potential role in FHB resistance. DON reportedly inhibited root growth when grown on agar medium, and functional DON detoxification of UGTs could reduce this inhibition (Ma et al., 2010; Pasquet et al., 2016). We overexpressed wheat TaUGT6 in Arabidopsis and found that the transgenic lines could increase root tolerance to DON. To date, few UGTs have been transformed into wheat. barley HvUGT13248 and its homolog Bradi5g03200 in B. distachyon were introduced into wheat by different research groups, and all of the data showed that expression of these genes increased FHB resistance and root tolerance to DON in wheat.
AUTHOR CONTRIBUTIONS

YH and HM conceived the experiments. YH, LW, XL, and LY performed most of the experiments and analyzed the data. PJ and XZ provided analytical tools and assisted in analyzing the data. JQ and GW performed toxin and HPLC-MS/MS analysis. XZ provided project administration. YH, HM, and XZ wrote the article. 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/fpls.2020.574775/full#supplementary-material

Supplementary Figure 1 | Alignment of the amino acid sequences of TaUGT6 and the UGTs previously reported to contribute to FHB resistance. Red box indicates the PSPG domain. TaUGTs were derived from Triticum aestivum, HvUGT1328 from Hordeum vulgare, DOGT1 from Arabidopsis thaliana, and Bradi5g03300 from Brachypodium distachyon.

Supplementary Figure 2 | Statistical analysis of the proportion of symptomatic spikelet between WT and the two T3 overexpressing transcript lines in the greenhouse in plastic pots (error bars indicate standard error and stars indicate significant differences; Student’s t-test, P < 0.05).

Supplementary Table 1 | List of primers used in this study.
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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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