A previously unknown zinc finger protein, DST, regulates drought and salt tolerance in rice via stomatal aperture control

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Abiotic stresses, such as drought and salinity, lead to crop growth damage and a decrease in crop yields. Stomata control CO₂ uptake and optimize water use efficiency, thereby playing crucial roles in abiotic stress tolerance. Hydrogen peroxide (H₂O₂) is an important signal molecule that induces stomatal closure. However, the molecular pathway that regulates the H₂O₂ level in guard cells remains largely unknown. Here, we clone and characterize DST (drought and salt tolerance)—a previously unknown zinc finger transcription factor that negatively regulates stomatal closure by direct modulation of genes related to H₂O₂ homeostasis—and identify a novel pathway for the signal transduction of DST-mediated H₂O₂-induced stomatal closure. Loss of DST function increases stomatal closure and reduces stomatal density, consequently resulting in enhanced drought and salt tolerance in rice. These findings provide an interesting insight into the mechanism of stomata-regulated abiotic stress tolerance, and an important genetic engineering approach for improving abiotic stress tolerance in crops.

Keywords: Rice; zinc finger protein; drought tolerance; salt tolerance; stomatal aperture; hydrogen peroxide

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Drought and salinity are two main abiotic stresses negatively affecting plant growth and seed production. Plants respond to and adapt to these stresses in order to survive. The understanding of plant responses to stresses in physiology, genetics, and molecular biology will be greatly helpful in improving the tolerance of plants to abiotic stresses through genetic engineering. Stomatal pores that are located in the epidermis of plant leaves control the uptake of CO₂ for photosynthesis and the water loss during transpiration, and play a crucial role in abiotic stress tolerance (Schroeder et al. 2001a; Hetherington and Woodward 2003). Under drought stress, abscisic acid (ABA)-mediated stomatal closure is a mechanism that plants use to adapt to water deficiency [Schroeder et al. 2001b; Zhu 2002; Fan et al. 2004]. Reactive oxygen species (ROS) including hydrogen peroxide (H₂O₂) that are widely generated under stress have been proposed to function as second messengers in ABA signaling in guard cells [McAinsh et al. 1996; Pei et al. 2000; Kohler et al. 2003; Bright et al. 2006]. Early work has shown that H₂O₂ can induce stomata closure [McAinsh et al. 1996; Zhang et al. 2001]. In guard cells, ABA-stimulated ROS accumulation activates plasma membrane calcium channels and triggers stomatal closure [Hamilton et al. 2000; Pei et al. 2000].

ABA-induced ROS are mainly generated by NADPH oxidase in guard cells [Kwak et al. 2003]. Mutation of two genes encoding subunits of an NADPH oxidase (AtrbohD and AtrbohF) impairs ABA-induced stomatal closing [Kwak et al. 2003]. To cope with increased levels of ROS, plants have evolved at least four kinds of ROS scavenging enzymes including catalase, superoxide dismutase, ascorbate peroxidase, and glutathione peroxidase. It has been shown that H₂O₂ is scavenged by glutathione peroxidase in the guard cells (Miao et al. 2006). So far, the genes that regulate the expression of H₂O₂ scavenging enzymes have not been identified, and the mechanism of regulation of H₂O₂ levels, especially in the stomata, is largely unknown.

Previous studies have shown that the transcriptional factors play important roles in plant tolerance to abiotic stresses [Stockinger et al. 1997; Liu et al. 1998; Haake et al. 2002; Yamaguchi-Shinozaki and Shinozaki 2006]. The Cys-2/His-2-type (C2H2) zinc finger protein—also called a classical or TFIIIA-type zinc finger—that was first found in Xenopus oocytes [Miller et al. 1985] is an important class of eukaryotic transcription factors. Several plant members of this family have various regulatory roles in stress responses and developmental processes.
However, it is still not known whether the C2H2-type zinc finger protein is involved in regulating stomatal aperture, although previous studies have shown that several transcription factors have been associated with stomata movement (Cominelli et al. 2005; Liang et al. 2005; Hu et al. 2006). In this study, we characterized a novel C2H2 zinc finger transcription factor, DST [drought and salt tolerance], that controls stomatal aperture under drought and salt stress in rice. DST contributes to stomata movement via regulation of genes involved in ROS homeostasis. These findings will shed light on the process of stomatal movement in plants and on the engineering of drought and salt tolerance in crops.

**Results**

**Isolation of dst mutant**

We performed a large-scale screen for the mutants to identify genetic loci that affect drought and salt tolerance in rice and isolated a dst mutant line. When treated with 20% PEG4000 (simulation of drought stress), the dst mutant exhibited less severe wilting than wild-type [ZH11] plants [Fig. 1A, top panel]. Furthermore, ~70% of dst mutant plants survived under 140 mM NaCl treatment, but nearly all of the wild-type plants wilted [Fig. 1A; Supplemental Fig. S1A]. The dst mutant exhibited better recovery than wild-type plants from both drought and salt stresses [Fig. 1A]. To accurately evaluate
the effect of the DST mutation on drought tolerance, we performed a soil drought experiment in deep polyvinyl chloride (PVC) pipes. The dst mutant plants showed stronger drought tolerance than the wild-type plants (Fig. 1A, bottom panel). These results indicated that mutation of the DST locus significantly improves drought and salt tolerance. There were no significant morphological alterations in dst mutants, except for a markedly wider leaf width [Fig. 1B,C]. Most agronomic traits were unchanged in the mutants, with the exception of the panicle number per plant and the main panicle length (Supplemental Fig. S2).

Physiological analysis of dst mutant
To elucidate the physiological mechanism of drought and salt tolerance in the dst mutant, we first investigated the water status in wild-type and dst mutants. Under dehydration stress, the dst mutant lost less water and maintained higher water content than wild-type plants [Fig. 1D,E]. The dst mutant also maintained higher osmolality during drought treatment [Fig. 1F]. These results indicate that the enhanced drought tolerance of the dst mutant is due to an increased ability to maintain water. Finally, we compared the K+ and Na+ content in the roots and shoots in response to NaCl stress. There was no significant difference in K+ content between the roots of the wild type and dst mutant; however, K+ content was lower in the shoots of dst mutants under both normal and salt stress conditions (Fig. 1G). NaCl treatment resulted in a lower accumulation of Na+ in the roots and shoots of dst mutant plants [Fig. 1G], indicating that the enhanced salt tolerance of dst mutant might be due to a reduction of Na+ toxicity. These results demonstrate that the DST mutation leads to enhancement of drought and salt tolerance in rice plants.

Stomatal movement changes in the dst mutant due to H2O2 accumulation
Because stomata are involved in responses to abiotic stresses (Hetherington and Woodward 2003), we examined the stomatal status of dst mutant and wild-type plants. The results showed that 42.4% of stomata were completely closed in the dst mutant plants, but only 16.7% were completely closed in the wild-type plants; also, only 21.2% of stomata were completely open in the dst mutants, but 50.8% were completely open in wild-type plants [Fig. 2A,B]. The percentage of partially open stomata was similar in dst mutant and wild-type plants (32.5%–36.4%). These results showed that stomatal movement is greatly affected in dst mutants. A significant decrease in stomatal density was also found in dst mutant plants [Fig. 2C], whereas there was no significant difference in guard cell size between dst mutant and wild-type plants [data not shown]. Meanwhile, the stomatal conductance was lower in dst mutant plants than that of wild-type plants [Fig. 2D]. We further investigated the stomatal status of the dst mutant under stress conditions, and found that the stomatal conductance of the dst mutant was significantly lower than that of the wild-type plants not only in the normal growth condition, but also in the salt stress condition [Supplemental Fig. S3]. These results demonstrate that the enhanced drought tolerance displayed by the dst mutant is primarily due to increased stomatal closure and decreased stomatal density, which prevent water loss, and that the enhanced salt tolerance of the dst mutant might be the result of a low transpiration rate that reduces Na+ transport from roots to shoots.

The phytohormone ABA induces stomatal closure (Schroeder et al. 2001b; Zhu 2002). However, measurement of endogenous ABA content revealed no significant difference between dst mutant and wild-type plants [Fig. 2E]. Therefore, the increased stomatal closure of the dst mutant is not the result of ABA accumulation. Since H2O2 induces leaf stomatal closure (McAinsh et al. 1996; Hamilton et al. 2000; Apel and Hirt 2004; Bright et al. 2006), we examined H2O2 content in rice leaves and found a higher accumulation in dst mutant plants [Fig. 2F,G]. Furthermore, by using a fluorescence dye, 2′,7′-dichlorodihydrofluorescein diacetate (H2DCFDA), we found that dst mutant plants accumulated more H2O2 in the guard cells than those of wild-type plants [Fig. 2H,I]. These results suggested that accumulation of H2O2 in guard cells probably underlies the increased stomatal closure in dst mutant plants.

DST encodes a novel zinc finger protein of unknown function
A map-based cloning approach was used to isolate the DST gene. Since the wide leaf is a typical and visible phenotype of the dst mutant [Fig. 1B], it was adopted as the standard dst phenotype. All F1 progeny showed a narrow leaf phenotype similar to that of wild-type plants [data not shown]. In the F2 population, narrow and wide leaf plants segregated in a 3:1 ratio [241 narrow vs. 63 wide; $\chi^2 = 2.96 < \chi^2_{0.05} = 3.84$; $P > 0.05$], demonstrating that the wide leaf phenotype of the dst mutant is controlled by a single recessive nuclear gene. The DST locus was initially mapped to the long arm of chromosome 3 between the markers H2423 and H2519 [Fig. 3A; Supplemental Table S1], and was subsequently fine-mapped to a 14-kb region between H2423 and H2437 [Fig. 3B]. Only one candidate gene, which encodes a 301-residue polypeptide with unknown function, localized in this region [Fig. 3C]. Protein sequence analysis showed that DST contains a nuclear localization signal (NLS) peptide and a single C2H2-type zinc finger motif at the N terminus [Fig. 3C,D]. A BLAST search found only one copy of DST in the rice genome. A comparison of the nucleotide sequences of the gene in dst mutant and wild-type plants revealed that two nucleotide substitutions lead to two amino acid substitutions: asparagine [N69] to aspartic acid [D] and alanine [A162] to threonine [T] [Fig. 3C], of which the N69 is a conserved residue of the zinc finger motif.

A complementation test was carried out to confirm the mapping result. Transfer of a 4.632-kb wild-type DNA fragment containing the DST promoter region and the entire ORF into dst mutants conferred phenotypes
similar to those of wild-type plants, including narrow leaf shape, sensitivity to drought and salt stresses, stomatal opening level, and H$_2$O$_2$ content (Fig. 3E; Supplemental Fig. S1A,B,D,E), whereas a knockout of DST using RNAi technology produced the dst phenotypes (Fig. 3E; Supplemental Fig. S1A–E). These results demonstrate that DST is indeed responsible for the phenotypic changes found in dst mutants, and that DST is a negative regulator of drought and salt tolerance.

We constructed N69D (M1) and A162T (M2) mutants to evaluate the role of the two substitutions (Fig. 3C). These two constructs and the dst (M, containing the two mutations) were each transformed into the dst mutant. Only those transformants expressing the M2 construct complemented dst phenotypes (Fig. 3F). Therefore, the mutation of N69, but not A162 is primarily responsible for the phenotypes of the dst mutant.

Subcellular localization and expression pattern of DST

Protein structure prediction showed that DST has a putative NLS [Fig. 3C]. We transiently expressed the DST-GFP fusion protein in onion epidermal cells and confirmed that the fusion protein specifically localizes to the nucleus [Fig. 4A]. The fluorescence of DST-GFP was also detected exclusively in the nucleus in transgenic rice plants that were transformed with a DST-GFP fusion construct under the control of the cauliflower mosaic virus (CaMV) 35S promoter (Supplemental Fig. S4). The tissue expression pattern of DST was determined by transforming rice plants with a β-glucuronidase (GUS) construct driven by the DST promoter. Strong GUS signals were detected in vascular tissues of the roots, internodes, and leaf sheath [Fig. 4B–H]. In the leaves, the GUS signal was also detected in the stomata apparatus.
To confirm the expression of DST in the stomata apparatus, we generated transgenic rice plants that were transformed with a DST-GFP fusion construct under the control of the DST native promoter. Confocal microscopic analysis showed that the fluorescence of DST-GFP was detected in the guard cells and the subsidiary cells (Fig. 4K; Supplemental Fig. S5), which was consistent with the role of DST in stomata regulation.

We also tested whether the expression of DST is regulated by drought and salt stresses. Real-time PCR analysis showed that DST is down-regulated rapidly after 0.5 h of drought or salt treatment, but that the expression resumed continuously after 24 h of treatment (Fig. 4L). These data are consistent with its role in regulating stomatal movement under drought and salt stresses. To exclude the possibility that DST is regulated by a circadian rhythm, we analyzed the rhythmic profiles of DST and found that DST is not regulated significantly by a circadian rhythm [Supplemental Fig. S6].

Figure 3. Map-based cloning of DST and complementation test. (A) The DST locus was mapped to chromosome 3 between markers H2423 and H2519 using 310 homozygous F2 plants. (B) Fine mapping of the DST locus. The DST locus was narrowed to a 14-kb region between markers H2423 and H2437. Numbers below the horizontal line in A and B are the number of recombinants. (C) The protein structure of DST. Two nucleotide substitutions in dst lead to an asparagine-to-aspartic acid change at amino acid 69 and an alanine-to-threonine change at amino acid 162. (D) Amino acid sequence alignments of the zinc finger domain. Identical and similar residues are displayed in colored boxes. The asterisk marks the cysteine and histidine in the C2H2-type zinc finger domain. The solid triangle denotes the point mutation in the zinc finger domain. GenBank accession numbers are ZFP1 (NP_178188), ZFP2 (NP_200560), ZFP5 (NP_172518), ZFP8 (NP_181725), SGT1-like protein (BAD73077), SUPERMAN (AAC49116), RAMOSA1 [AAA71040], and LIF [BAB58897]. (E) Drought and salt treatment of T2 transgenic lines including vector control, complementation, and RNAi. Twenty-day-old plants [top] were treated with 18% PEG4000 for 10 d [middle] or 100 mM NaCl for 15 d [bottom]. (F) Drought and salt treatment of T2 transgenic lines with point mutation constructs. Twenty-day-old plants [top] were treated with 18% PEG4000 for 15 d [middle] or 100 mM NaCl for 20 d [bottom]. M1, M2, and M indicate transgenic plants with N69D, A162T, and both point mutation constructs, respectively.

DST is a transcription factor with transactivational activity

We performed a transcriptional activation assay and found that expression of the DST-BD [GAL4-binding domain] fusion protein in yeast resulted in high reporter gene expression [Fig. 5A], which revealed that DST has strong activity as a transcription activator. Assessment of the two amino acid substitutions in the dst mutant showed that the N69D substitution decreased the level of transcriptional activation, while the A162T substitution did not alter the activation level. These results indicated that the N69 of DST is required for transcriptional activation. Interestingly, when the two amino acid substitutions were combined, activation was lower than that of the N69D mutation alone [Fig. 5A], possibly because the two amino acid substitutions modify DST protein configuration. Further quantitative assays of the β-galactosidase activity confirmed these results [Supplemental Fig. S7]. Deletion analysis shows that the first 72 amino acids of DST contribute to its transcriptional activation activity.
acids, containing the zinc finger motif, are required for transcriptional activation, and that the activation domain is localized in the first 145 N-terminal residues [Fig. 5A].

DST binds to the TGCTANNATTG element by the zinc finger domain

We used a bacterial one-hybrid system [Meng et al. 2005] to identify the DNA-binding sequence of DST. Among the sequences identified, six sequences had a conserved TGNTANN(T/A)T sequence [Fig. 5B]. Electrophoretic mobility shift assay [EMSA] showed that the His-DST fusion protein, but not His alone, could bind to a B3 sequence that contains this conserved sequence, whereas an excess unlabeled B3 sequence effectively reduced or abolished the His-DST binding to the labeled B3 sequence [Fig. 5C]. Meanwhile, the His-DST protein could bind to the B3 sequence in a dosage-dependent manner [Supplemental Fig. S8A]. These results demonstrated that DST binds specifically to this sequence. Individual mutations of TG [M10], TA [M3], and AT [M5], and simultaneous mutations [M11 and M12] in the TGNTANNAT sequence markedly reduced or abolished the binding of His-DST [Fig. 5D,E]. The mutants [M4, M7, M8, M9, and M13] with the conserved nucleotides did not significantly reduce the binding of His-DST (Fig. 5D–F). However, other mutants [M6 and M14] markedly reduced the binding of His-DST, showing that TG and C are also required for DST binding [Fig. 5D–F]. Taken together, these results showed that DST binds to the TGCTANNATTG sequence, which is a novel cis-element and named DBS [DST-binding sequence]. This was further demonstrated by those mutations (M15, M16, and M17) in this core sequence that completely abolished the binding of His-DST [Fig. 5D–F]. Interestingly, the M8 mutant, which produced new ATTG core nucleotides, increased the binding of His-DST (Fig. 5E; Supplemental Table S2).

We also tested whether the mutant DST protein in dst mutants could bind to DBS, and found that the mutant form of DST protein could bind to DBS as the wild-type protein did (Supplemental Fig. S8B–D).

Based on previous studies [Wolfe et al. 2000; Sakamoto et al. 2004; Lin et al. 2007], we assumed that the zinc finger motif of DST might be the DNA-binding domain. The preincubation of the His-DST fusion protein with two metal chelators—EDTA or 1,10-phenanthroline—effectively reduced or abolished the DNA-binding ability of His-DST, suggesting that DST binds to the DBS element by the N-terminal zinc finger motif [Fig. 5G]. This suggestion was further confirmed by EMSA using a truncate DST protein with deletion of the zinc finger domain (Supplemental Fig. S8B–D).

DST targets to genes related to H2O2 homeostasis

Expression profile analysis using the Affymetrix Rice Genome Genechip was used to identify DST target genes. We found that among ∼50,000 rice genes, the transcription levels of 60 genes were altered significantly [more than threefold change] in dst mutant plants [Supplemental Table S3]. A high percentage (60%) of these 60 genes had lower expression levels in dst mutants (Supplemental Table S3), and mutation of the DST protein in dst mutants lost its transactivation activity [Fig. 5A], indicating that DST might promote expression of these genes. It is
noteworthy that at least five of the down-regulated genes encode proteins related to ROS homeostasis: two peroxidases (24 precursor, 12 precursor; EC1.11.1.7), glutathione S-transferase (OsGSTU2; EC2.5.1.18), and two cytochrome P450s (71D10, 94A2; EC1.14.-.-) (Jakoby 1978; Marrs 1996; Noctor and Foyer 1998; Asada 1999; Rodriguez Milla et al. 2003; Mittler et al. 2004, 2005; Campos et al. 2005; Moller et al. 2007). The expression of these genes was greatly down-regulated in dst mutant plants (Supplemental Table S3), four of which were confirmed by RT–PCR (Supplemental Fig. S9).

We subsequently searched the promoter regions of four genes and found two or three DBSs in each promoter (Supplemental Fig. S10). The promoter sequences of two genes were used for EMSA, showing that DST can bind to the promoter sequences (Fig. 6A). Furthermore, a chromatin immunoprecipitation (ChIP) assay using wild-type plants showed that the anti-DST antibody specifically precipitated promoter sequences of all four genes from the cells of wild-type plants (Fig. 6B), indicating that DST binds directly to the promoters of these genes in vivo. Noticeably, in these five genes, peroxidase 24 precursor may directly scavenge H2O2. Peroxidase activity assay confirmed that peroxidase 24 precursor catalyzes the reduction of H2O2 (Fig. 6C,D). Real-time PCR analysis showed that the expression level of peroxidase 24 precursor was higher in the guard cell protoplast (GCP)-enriched sample compared with the mesophyll cell.

**Figure 5.** DST has transactivation activity and DNA-binding functions. (A) Schematic diagrams of various constructs used for transactivation activity assay. (DNA-BD) GAL4 DNA-binding domain; (GAL4 AD) GAL4 activation domain; (NLS) NLS sequence; (MCS) multiple cloning site; (CDS) coding sequence of DST. (B) Alignment of representative sequences identified in bacterial one-hybrid experiment. Conserved nucleotides are shown in color. Sequence logo was generated using Weblogo [see the Materials and Methods for detail]. (C) EMSA showing that the His-DST fusion protein, not His alone, can bind the B3 sequence. Competitive binding analysis was performed with increasing amounts of unlabeled probe. The amount of competitive unlabeled probe in lanes 4–7 was 10, 50, 100, and 200 times of that of the labeled probe, respectively. (D) B3 sequence and its various mutants. (E,F) EMSA with the same amount of wild-type B3 probe or mutant probes. (G) Two metal chelators, EDTA or 1,10-phenanthroline, effectively abolish the binding of DST to B3 probe, suggesting that the zinc finger motif of DST might be the DNA-binding domain. The arrow, asterisk, and triangle in C and E–G indicate the up-shifted bands, nonspecific binding, and free probe, respectively. (FP) Free protein.
protoplast (MCP)-enriched sample (Fig. 6E). This result indicated that peroxidase 24 precursor was expressed in guard cells, which was consistent with the expression of DST in the guard cells. Furthermore, peroxidase 24 precursor was not only down-regulated in the leaf, but also in the GCP of the dst mutant (Fig. 6F). Meanwhile, under salt and drought treatment, peroxidase 24 precursor was down-regulated and then resumed (Fig. 6G), which was similar to the expression pattern of DST. These results indicated that DST might directly regulate the expression of peroxidase 24 precursor under salt and drought stresses. Collectively, we propose that accumulation of H$_2$O$_2$ in the dst mutant (Fig. 2F–I) may be the result of down-regulation of the genes related to H$_2$O$_2$ homeostasis.

**Discussion**

Generally, C2H2-type zinc finger proteins contain more than one zinc finger motif for DNA binding in the N terminus and separable transcriptional activation (or repression) domains in the C terminus (Wolfe et al. 2000). The zinc finger motif is thought to recognize and bind its target DNA sequence, but has never been shown to be required for transcriptional activity (Wolfe et al. 2000; Sakamoto et al. 2004; Lin et al. 2007). However, here we...
surprisingly found that the single zinc finger motif of DST is required not only for its DNA-binding activity but also its transcriptional activity, indicating that DST is a novel transcription factor. It brings us brand-new knowledge and a new understanding about the zinc finger transcription factor.

H$_2$O$_2$ plays an important role in mediating ABA-induced stomatal closure [McAinsh et al. 1996; Pei et al. 2000; Kohler et al. 2003; Bright et al. 2006]. In Arabidopsis guard cells, H$_2$O$_2$ was scavenged by glutathione peroxidase [Miao et al. 2006]. Mutation of Arabidopsis thaliana glutathione peroxidase 3 (ATGPX3) leads to the accumulation of H$_2$O$_2$ in guard cells. However, direct upstream regulators of genes related to H$_2$O$_2$ homeostasis, such as ATGPX3, have not been identified. Here, we showed that rice peroxidase 24 precursor has peroxidase activity to scavenge H$_2$O$_2$ [Fig. 6C,D]. Meanwhile, EMSA and ChIP results showed that DST targets to the promoter sequence of peroxidase 24 precursor [Fig. 6A,B]. Therefore, DST is, to our knowledge, the first known direct upstream regulator of the genes related to H$_2$O$_2$ homeostasis in guard cells. Mutation of DST results in the down-regulation of peroxidase 24 precursor [a scavenger of H$_2$O$_2$], might lead to the accumulation of H$_2$O$_2$ in guard cells and trigger stomatal closure, and enhances drought and salt tolerance. Recently, Arabidopsis mutants lacking either or both a cytosolic and chloroplastic ascorbate peroxidase, which were responsible for H$_2$O$_2$ removal, were found to be more tolerant to salinity stress [Miller et al. 2007]. Interestingly, two ROS scavengers [glutathione S-transferase GSTU6 and peroxidase 16 precursor] were found to be up-regulated in the dst mutant [Supplemental Table S3], and they might contribute to the removal of excess ROS because high levels of ROS have been shown to be toxic [Apel and Hirt 2004].

In the last decade, many investigators have made progress in illuminating the regulatory pathways involved in abiotic stress tolerance [Zhu 2002; Shinozaki et al. 2003; Yamaguchi-Shinozaki and Shinozaki 2006]. For example, dehydration-responsive element (DRE)/DRE-binding protein (DREB) pathways [Stockinger et al. 1997; Liu et al. 1998]; the ABA-responsive element (ABRE)/ABRE-binding protein ABI5 pathway [Choi et al. 2000; Finkelstein and Lynch 2000; Lopez-Molina and Chua 2000; Uno et al. 2000]; and the MYB recognition sequence (MYBRS)/MYB transcription factor pathway [Abe et al. 1997, 2003]. The cis-acting and trans-acting regulatory elements in these pathways have been precisely analyzed at the molecular level [Yamaguchi-Shinozaki and Shinozaki 2005]. DBS is a novel cis-acting regulatory element that was found in the promoter sequences of the genes related to ROS homeostasis. DST binds to DBS and regulates the expression of ROS homeostasis-related genes. In the initial stage of dehydration stress, the expression of DST was repressed, resulting in the down-regulation of peroxidase 24 precursor—whose promoter contains DBS—giving rise to H$_2$O$_2$ accumulation and promoting stomatal closure.

ABA is an important regulatory component of plant responses to abiotic stresses such as drought and salinity. Many genes that are induced by drought and salt treatment also are induced, or not, in response to exogenous ABA treatment, suggesting that there are ABA-dependent or ABA-independent stress response pathways [Zhu 2002; Yamaguchi-Shinozaki and Shinozaki 2006]. Treatment of exogenous ABA did not significantly affect the expression of DST, and there were no significant differences between stomatal openings among wild type, the dst mutant, the complementation line, and the RNAi line under ABA treatment [Supplemental Fig. S11A,B], indicating that the pathway that DST was involved in was ABA-independent.

Guard cells or stomatal apparatus cells are important cell types for dehydration tolerance. Although the transcriptional network in these cells is quite important in understanding plant response mechanisms to dehydration and for improvement of dehydration tolerance of crops, progress in this area has been relatively limited. The DST regulatory pathway revealed in this study will lead to better understanding of the transcription network in the stomatal apparatus. In addition, DST also has pleiotropic effects—at least on stomatal density and leaf width—implying that DST also is involved in regulating the development of tissues or organs. How DST regulates stomatal density and leaf width is an intriguing question to be answered in future studies. Based on our data, we summarized a model to explain the potential role of DST in the regulation of stomatal status and abiotic stress tolerance [Fig. 7].

Engineering of the stomatal response to reduce water loss would be a powerful tool for the enhancement of drought tolerance in crops [Schroeder et al. 2001b]. Down-regulation of DST function, which constitutively reduces the stomatal opening and stomatal density, would allow

Figure 7. A proposed model for the role of DST in regulation of abiotic stress tolerance. DST binds directly to the DBS element in the promoter of genes related to H$_2$O$_2$ homeostasis and activates their transcription, thereby inhibiting H$_2$O$_2$ accumulation. Inhibition of H$_2$O$_2$ accumulation influences stomatal closure and, ultimately, abiotic stress tolerance. However, the pathway for regulation of stomatal density remains to be elucidated in future studies. Under drought and salt stress, the expression of DST was repressed, resulting in the down-regulation of genes related to H$_2$O$_2$ homeostasis, such as peroxidase 24 precursor, whose promoter contains DBS, giving rise to H$_2$O$_2$ accumulation and promoting stomatal closure, and finally enhancing drought and salt tolerance. This pathway is ABA-independent, and is different from the ABA-induced H$_2$O$_2$ accumulation pathway in controlling stomatal closure.
engineering of crop plants [such as rice] to reduce water loss, enhance drought tolerance, and reduce water requirements in agricultural regions with marginal fresh water availability. Moreover, the down-regulation of DST does not affect rice grain yield (Supplemental Fig. S2F), so although the DST mutation results in a smaller stomatal opening and a lower stomatal density [Fig. 2A–C], it is sufficient for CO₂ influx and growth. This characteristic shows the strong potential value of DST in abiotic stress tolerance engineering. Therefore, our findings not only provide new insights into the molecular mechanisms underlying abiotic stress tolerance in plants, but also facilitate molecular breeding efforts to improve drought and salt tolerance in staple crops and the conservation of water resources in agricultural regions where fresh water is scarce.

Materials and methods

Plant material and growth conditions

The wild-type plant Zhonghua 11 (ZH11) is a drought- and salt-sensitive *japonica* variety. We screened >270,000 ethyl methanesulfonate (EMS)-mutagenized M2 rice seedlings (~9000 lines) treated with 140 mM NaCl. In the first round of screening, >10 lines were isolated as putative salt tolerance mutant plants. Among these mutant lines, one mutant maintained the highest ability to survive in the treatment of NaCl. When treated with 20% PEG4000 (similation of drought stress), this mutant exhibited less severe wilting than wild-type (ZH11) plants. This mutant was named as dst. The dst mutant was backcrossed to the wild-type and then a dst mutant line after propagation for at least five generations was used for further analysis. Seeds were placed for 1 wk or more at 42°C to break any possible dormancy, soaked in the water at room temperature for 3 d, and then germinated for 1 d at 37°C. The most uniformly germinated seeds were sown in a 96-well plate from which the bottom was removed. The plate was floated on water for 1 d at 37°C in the dark to encourage root growth, before removal to a growth chamber with a 13-h light (24°C)/11-h dark (20°C) photoperiod. Five days later, the seedlings were cultured with Yoshida's culture solution with or without 1% hygromycin (for the selection of transgenic plants). For salt treatment, 20-d-old seedlings were transferred to culture solution containing 100 mM or 140 mM NaCl. For PEG treatment, 20-d-old seedlings were transferred to culture solution containing 18.85% or 20% (w/v) PEG4000. For soil drought experiments in PVC pipes, 25-d-old seedlings cultured in the growth chamber were transplanted to PVC pipes and cultured in the greenhouse at 24°C–30°C and 50%–60% relative humidity. Thirty days after transplantation, water was removed from the bottom of the PVC pipes for dehydration treatment.

**Diaminobenzidine (DAB) staining and quantitative measurement of H₂O₂**

H₂O₂ was detected by DAB staining as described previously (ThordalChristensen et al. 1997). The fully expanded leaves of 20-d-old wild type and dst plants were detached and vacuum-infiltrated with the DAB solution. The sampled leaves were placed in a growth chamber for 8 h at 28°C and cleared in boiling ethanol (96%) for 10 min before photographing. A measurement of H₂O₂ production was performed by extracting H₂O₂ from leaves according to a previously described method (Rao et al. 2000) and quantifying the molecule with an Amplex Red Hydrogen/Peroxidase Assay Kit (Molecular Probes) according to the manufacturer's instructions.

**Detection of H₂O₂ production in guard cells**

H₂O₂ production in guard cells was detected using H₂DCFDA (Molecular Probes) as described previously (Zhang et al. 2001) with modifications. Leaves of 45- to 50-d rice seedlings were immersed in 0.01% Tween-20 and vacuum-infiltrated for 5 min. After rinsing twice with distilled water, leaves were incubated in 2% (w/v) cellulase RS [Yakult Honsha] for 5 h at 40°C without shaking to facilitate peeling off the epidermal strips. The epidermal strips were peeled off from leaves using tissue forceps. After washing with loading buffer [10 mM Tris-HCl, 50 mM KCl at pH 7.2], the epidermal strips were incubated in staining buffer (loading buffer containing 50 μM H₂DCFDA) for 10 min at room temperature in the dark. The epidermal strips were washed with distilled water to remove the excess dye. The fluorescence was examined using a confocal laser-scanning microscope [Carl Zeiss, LAM510] with the following settings: excitation, 488 nm; emission, 525 nm; and power, 5%. All confocal images were taken under identical conditions: 12 bits; scanning speed, 7; frame, 1024 × 1024; pinhole, 300, and detector gain, 750. For quantifying the gray value (0–4095) in the guard cell, the guard cell region was selected, and the mean gray value was recorded. For dst and wild-type plants, nine independent experiments were performed, and >10 stoma were selected randomly for analysis in each experiment.

**Imaging of rice stomata and measurement of stomatal conductance**

Leaves of 25-d-old plants were detached and immediately fixed by liquid nitrogen. The stomatal pictures were obtained using an environmental scanning electron microscopy (XT30 ESEM-TMP, Philips). For the stomatal conductance measurement, seedlings were cultured in a growth chamber for 25 d, and then transplanted to soil in a greenhouse under natural growth conditions. At tillering stage (~35 d after transplant), the second fully expanded leaf, counting from the top of plant, was used for stomatal conductance measurement with a portable gas analysis system (LI-COR 6400, LI-COR, Inc.). The measurement was performed under the following conditions: a constant water concentration of 18.85 ± 0.02 mmol mol⁻¹, a constant temperature of 28.80°C ± 0.02°C, and a constant CO₂ concentration of 350.20 ± 0.02 μL L⁻¹.

**Bacterial one-hybrid analysis and EMSA**

Bacterial one-hybrid analysis was performed as described previously (Meng et al. 2005) with some modifications. The full-length coding sequence of DST was PCR-amplified with primers containing BstXI and BamHI sites, and then cloned into the AvrII-BamHI sites of pPH1 (Supplemental Table S2). High-efficiency electrocompetent DH10B was substituted for XL1-blue. The 3-amino-1,2,4-triazole (3-AT) selection plate contained 3 mM or 4 mM 3-AT (Sigma). Twenty clones grown on 3 mM or 4 mM 3-AT selection plates were selected randomly and sequenced. Fifteen unique sequences were analyzed using the Gibbs Motif Sampler [http://bayesweb.wadsworth.org/cgi-bin/gibbs.12.pl?data_type=DNA]. Among them, six sequences containing a conserved sequence were identified. The sequence logo was generated using Weblogo 3 [http://weblogo.berkeley.edu/logo.cgi]. The plasmids pBH1 (Addgene plasmid 12611) and pH3U3-mcs [Addgene plasmid 12609] and bacteria strain US0his8-pyrF [Addgene plasmid 12614] were obtained from Professor Scot Wolfe through Addgene [http://www.addgene.org].
To generate the 6xHis-PRX, the ORF of activity assay
Preparation of 6xHis-PRX fusion protein and proxidase
DST fusion protein was prepared as described above and purified
HindIII–SalI site of pMAL-c2X (New England Biolabs). The MBP-
into the bacterial strain BL21 (DE3) pLysS. The transformed cells
HindIII, and inserted into the EcoRI–HindIII site of pET-32a
to release the ORF of DST
constructed by digesting the pDST plasmid with HindIII and SalI
using MBP-DST fusion protein. The MBP-DST plasmid was
purified with Ni-NTA resin (Qiagen).

Using a high-pressure cell crusher and the fusion proteins were
lysed by using a high-pressure cell crusher and the fusion proteins were purified with Ni-NTA resin (Qiagen) according to the manufacturer’s instructions.

The peroxidase activity assay was performed using an Amplex Red Hydrogen Peroxide/Peroxidase Assay Kit [Molecular Probes] according to the manufacturer's instructions. About 1 µg of 6xHis-PRX fusion protein was incubated with Amplex Red reaction mixture at 30°C. Absorbance at 560 nm was measured in a SpectraMax 190 [Molecular Devices] every 1 min for 60 min. Values were subtracted to the background absorbance. About 2 µg of 6xHis protein and 100 µU/mL horseradish peroxidase (HRP) were used as negative and positive controls. For generation of a standard curve, 0, 25, 37.5, 50, 62.5, 75, 87.5, and 100 µU/mL HRP were incubated with Amplex Red reaction mixture for 60 min at 30°C. After subtracting to the background absorbance, the values were used to generate a linear regression equation. The relative activity of 6xHis-PRX and 6xHis protein was calculated with equal mole protein according to the linear regression equation.

Other experimental methods are described in the Supplemental Material.

Accession numbers
Raw microarray data have been deposited in Gene Expression Omnibus (GEO, http://www.ncbi.nlm.nih.gov/geo) with GEO Series accession number GSE11175. The GenBank accession number for DST sequences is GQ178286.

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