

# A Wheat *SIMILAR TO RCD-ONE* Gene Enhances Seedling Growth and Abiotic Stress Resistance by Modulating Redox Homeostasis and Maintaining Genomic Integrity<sup>CJW</sup>

Shuantao Liu,<sup>1,2</sup> Shuwei Liu,<sup>2</sup> Mei Wang,<sup>2</sup> Tiandi Wei, Chen Meng, Meng Wang, and Guangmin Xia<sup>3</sup>

Key Laboratory of Plant Cell Engineering and Germplasm Innovation, Ministry of Education, School of Life Science, Shandong University, Jinan 250100, China

Plant growth inhibition is a common response to salinity. Under saline conditions, Shanrong No. 3 (SR3), a bread wheat (*Triticum aestivum*) introgression line, performs better than its parent wheat variety Jinan 177 (JN177) with respect to both seedling growth and abiotic stress tolerance. Furthermore, the endogenous reactive oxygen species (ROS) was also elevated in SR3 relative to JN177. The SR3 allele of *sro1*, a gene encoding a poly(ADP ribose) polymerase (PARP) domain protein, was identified to be crucial for both aspects of its superior performance. Unlike RADICAL-INDUCED CELL DEATH1 and other *Arabidopsis thaliana* SIMILAR TO RCD-ONE (SRO) proteins, *sro1* has PARP activity. Both the overexpression of *Ta-sro1* in wheat and its heterologous expression in *Arabidopsis* promote the accumulation of ROS, mainly by enhancing the activity of NADPH oxidase and the expression of NAD(P)H dehydrogenase, in conjunction with the suppression of alternative oxidase expression. Moreover, it promotes the activity of ascorbate-GSH cycle enzymes and GSH peroxidase cycle enzymes, which regulate ROS content and cellular redox homeostasis. *sro1* is also found to be involved in the maintenance of genomic integrity. We show here that the wheat SRO has PARP activity; such activity could be manipulated to improve the growth of seedlings exposed to salinity stress by modulating redox homeostasis and maintaining genomic stability.

## INTRODUCTION

Genetic analysis of the stress response of model plants has identified a number of key genes involved (Shinozaki and Yamaguchi-Shinozaki, 2000; Zhu, 2002; Munns and Tester, 2008), but few of these genes have as yet been genetically manipulated for crop improvement. Most attempts to enhance stress tolerance using a transgenic approach have failed because the transgene has typically exerted a negative effect on plant growth (Skirycz et al., 2011). Thus, to identify genes involved in both stress tolerance and growth improvement is important for crop breeding.

Salinity, in common with several other agents of abiotic stress, induces the production of toxic reactive oxygen species (ROS), which if not controlled, can ultimately trigger cell death (Dat et al., 2000). However, relatively low levels of ROS are implicated in stress tolerance. There is a substantial degree of support for the suggestion that maintaining a low level of endogenous H<sub>2</sub>O<sub>2</sub> enhances tolerance to a number of abiotic stresses (Mittler, 2002; Neill et al., 2002; Apel and Hirt, 2004). On the other hand, some reports support the idea that constitutive elevation of H<sub>2</sub>O<sub>2</sub>

in plants can improve multistress tolerance in plants (Van Breusegem et al., 2008; Huang et al., 2009). However, whether elevated ROS content can simultaneously increase both plant growth and abiotic stress tolerance is unknown.

The enzyme poly(ADP ribose) polymerase (PARP) is part of the early response to DNA breakage caused by radiative and oxidative stress (Briggs and Bent, 2011). PARP-like genes are present in many eukaryotes, and the PARP catalytic domain is characteristically associated with a range of other domains (Amé et al., 2004; Jaspers et al., 2010). A prominent example is represented by the *Arabidopsis thaliana* gene RADICAL-INDUCED CELL DEATH1 (*RCD1*), which contains a WWE domain and an RST (for RCD-SRO-TAF4) domain, both of which are responsible for ensuring the protein-protein interactions, and its expression in *Saccharomyces cerevisiae* strain WYT could overcome the oxidative stress sensitive phenotype of these yeast cells (Belles-Boix et al., 2000; Aravind, 2001). The combination of a PARP and an RST domain is specific to plants, and proteins carrying both have been termed SIMILAR TO RCD-ONE (SRO) (Jaspers et al., 2010). A loss-of-function mutation in *At-RCD1* results in an enhanced sensitivity to salt, Glc, and apoplastic ROS but increased resistance to freezing and chloroplastic superoxide formation by methyl viologen (MV) (Overmyer et al., 2000; Ahlfors et al., 2004; Fujibe et al., 2004; Katiyar-Agarwal et al., 2006; Teotia and Lamb, 2009). Double mutants in both *RCD1* and *SRO1* have severe developmental defects and behave differently in several developmental events and abiotic stress responses (Jaspers et al., 2009; Teotia and Lamb, 2009), which are similar to those seen in the stress-induced morphological response (Teotia et al., 2010), which is known to be associated with changes in redox balance (Potters et al., 2009). The *Arabidopsis* genome encodes at least

<sup>1</sup> Current address: Institute of Vegetables, Shandong Academy of Agricultural Sciences, Jinan 250100, China.

<sup>2</sup> These authors contributed equally to this work

<sup>3</sup> Address correspondence to xiagm@sdu.edu.cn.

The author responsible for distribution of materials integral to the findings presented in this article in accordance with the policy described in the Instructions for Authors (www.plantcell.org) is: Guangmin Xia (xiagm@sdu.edu.cn).

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www.plantcell.org/cgi/doi/10.1105/tpc.113.118687

three putative PARPs and six SROs. The former have high levels of sequence similarity with mammalian PARPs at the peptide level, but not all PARPs have poly(ADP ribosyl)ation activity; some show mono(ADP ribose) transferase (mART) activity and others appear to be enzymatically inactive (Citarelli et al., 2010). Although the ability to bind NAD<sup>+</sup> is necessary for the activity of PARP, At-RCD1 does not readily bind NAD<sup>+</sup> and has no detectable poly(ADP ribosyl)ation activity and mART activity (Jaspers et al., 2010). Other *Arabidopsis* SROs and rice (*Oryza sativa*) SRO1c were predicted to lack poly(ADP ribosyl)ation activity based on bioinformatics (Jaspers et al., 2010; You et al., 2013), and the *Arabidopsis* SROs were also reported to lack mART activity (Wang et al., 2011). Whether SROs in other plants are enzymatically active remains to be elucidated.

The salinity-tolerant bread wheat (*Triticum aestivum*) cultivar Shanrong No. 3 (SR3) is a derivative of an asymmetric somatic hybrid between bread wheat and tall wheatgrass (*Thinopyrum ponticum*) (Xia et al., 2003). This novel cultivar has been shown to be higher yielding than its wheat parent Jinan 177 (JN177) when grown in saline soil and has been planted commercially in a number of areas where the soil suffers from alkaline salinity (Xia, 2009). The genetic basis of much of its salinity tolerance has been shown to rely on ROS homeostasis based on the transcriptomic and proteomic outcomes in response to salinity stress (Peng et al., 2009; Liu et al., 2012). Here, we report the superior traits of SR3 and demonstrate that a candidate gene, which is the allele of the parent JN177 via point mutations, encodes an SRO protein containing a N-terminal WWE domain, a PARP-like domain, and a C-terminal RST domain. We found that the wheat SRO has PARP activity and DNA binding ability and is involved in redox homeostasis regulation and genomic stability maintenance. Our findings demonstrated that *sro1* could be effectively used to improve plant growth and stress tolerance.

## RESULTS

### The Higher Salinity Tolerance of Cultivar SR3

Compared with JN177, SR3 seedlings were larger in the normal field and kept their growth advantage under saline field conditions. More seedlings became yellow, wilted, and died in JN177 than in SR3 in the 0.5% salinity fields. To investigate SR3's abiotic stress responses more efficiently, we further investigated its phenotypes of growth and stress resistance at the seedling stage grown in culture solution. SR3 seedlings grew more vigorously than those of its bread wheat parent cultivar JN177, under both nonstressed and stressed (polyethylene glycol [PEG], NaCl, or H<sub>2</sub>O<sub>2</sub>) conditions (Figures 1A to 1D and 1I; Supplemental Figure 1A). Alterations in cellular ROS levels are known to affect the growth of plants, especially the root system (Finkemeier et al., 2005; Olmos et al., 2006; Miller et al., 2007). Therefore, we examined the ROS content in the roots of SR3 and JN177 under both nonstressed and stressed (PEG, NaCl, or H<sub>2</sub>O<sub>2</sub>) conditions. Consistently, the ROS content of SR3 roots was higher than that of JN177, irrespective of the treatment (Figures 1E to 1H and 1J; Supplemental Figure 1B) and was positively correlated with root length (Figure 1I), implying that the gene(s) responsible for ROS accumulation in SR3

is also responsible for the vigorous root growth under stressed conditions.

### Isolation of a Stress-Responsive Gene Belonging to the SRO Family

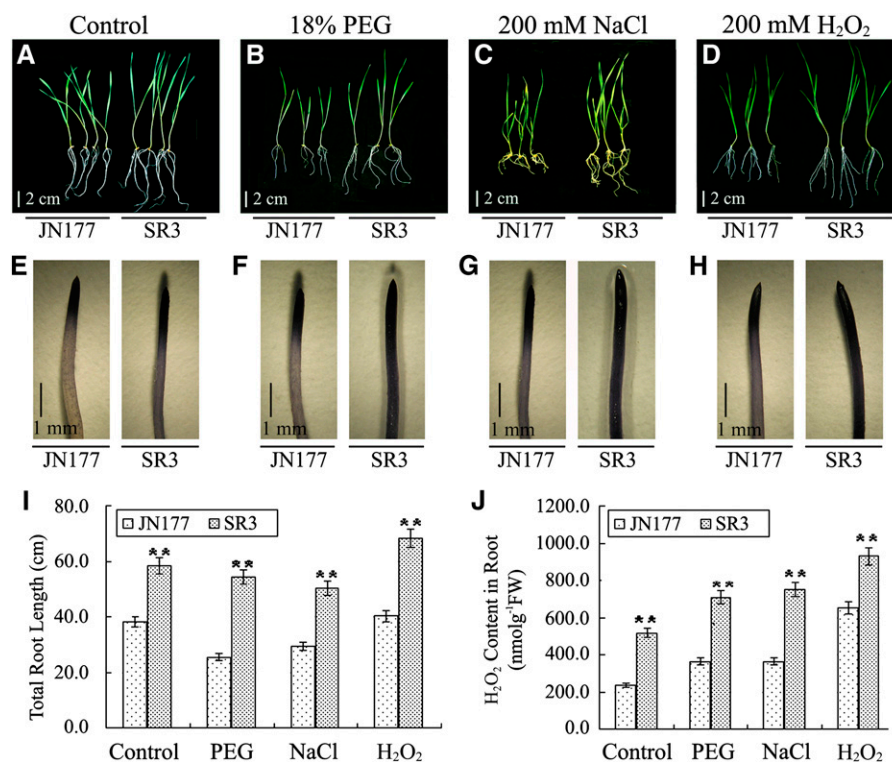
A set of 226 genes that were differentially expressed (by at least 2-fold) in SR3 seedlings under stress (200 mM NaCl or 18% PEG) was identified by microarray analysis (Liu et al., 2012). Out of these genes, those related to ROS homeostasis are suggested to play an important role in the generation of the above salinity-tolerant phenotype of cultivar SR3. Among the 64 genes related to oxidative stress (Supplemental Table 1), one shared a substantial degree of homology with At-RCD1. Copies of this gene have been located on the homoeologous group 5 chromosomes (<http://wheat.pw.usda.gov/GG2/blast.shtml>), which is also the site of the major stress tolerance locus in SR3 (our unpublished data). Expression analysis of this gene, based on quantitative RT-PCR, showed that both salinity and H<sub>2</sub>O<sub>2</sub> stress induced its expression within 1 h, earlier than the dehydration stress (PEG), and its expression remained upregulated for at least 24 h under all three treatments. Transcript abundance was consistently higher in the roots of SR3 than in those of JN177 (Figures 2A to 2C).

The full-length cDNA sequences of both the SR3 and JN177 alleles comprised a 1734-bp open reading frame, encoding a 578-residue 63.8-kD protein. The nucleotide sequences differed from one another at three positions (Gly-585, Gly-749, and Gly-1027 in JN177 to Ala-585, Thr-749, and Ala-1027 in SR3), resulting in two residues being converted from Gly-250 and Ala-343 in JN177 to Val-250 and Thr-343 in SR3 (Supplemental Figure 2). The nucleotide sequence of its closest rice putative ortholog Os-SRO1b was 75% similar, while the *Arabidopsis* genes At-RCD1 and At-SRO1 (involved in the response to oxidative and osmotic stress; Teotia and Lamb, 2009) were 47% similar (Figure 2D). The gene product included both an RST and a PARP-like domain at its C terminus and a WWE domain at its N terminus (Figure 2E) and thus belongs to the SRO protein family. By comparing all other SRO genes from SR3, JN177, and tall wheatgrass, the two genes show the closest relationship, which indicates that the SR3 allele is derived from the JN177 SRO1 via point mutation; therefore, we designated JN177 SRO1 Ta-SRO1 and the SR3 allele Ta-sro1.

To determine where Ta-sro1 functions in cells, we transiently expressed Ta-sro1 with a green fluorescent protein (GFP) tag under the control of a constitutive cauliflower mosaic virus 35S promoter in *Arabidopsis* mesophyll protoplasts. In contrast with the GFP control that diffused in both the nucleus and cytoplasm, the Ta-sro1-GFP fusion protein exclusively localized to the nucleus (Figure 2F).

### Constitutive Expression and Suppression of Ta-sro1 in Wheat

To determine if Ta-sro1 is the crucial gene that enhances both seedling growth and abiotic stress resistance in SR3, the function of Ta-sro1 was extensively explored both by constitutively expressing it in the salinity sensitive wheat cultivar Yangmai 11 and by abolishing its expression in SR3 (Supplemental Figure 3). As both wild-type Yangmai 11 and the SR3 Ta-sro1 RNA interference (RNAi) lines wilted rapidly when challenged with moderate levels



**Figure 1.** The Abiotic Stress Tolerance of Cultivar SR3.

(A) to (D) Phenotype of 1-week-old JN177 and SR3 under control (A), 18% PEG (B), 200 mM NaCl (C), or 200 mM H<sub>2</sub>O<sub>2</sub> (D). Nonstressed or stressed seedlings grown for 2 weeks. JN177 is a bread wheat cultivar. The salinity tolerant bread wheat cultivar SR3 is a derivative of an asymmetric somatic hybrid between JN177 and tall wheatgrass.

(E) to (H) Root tips of 1-week-old JN177 and SR3 seedlings under control (E), 18% PEG (F), 200 mM NaCl (G), or 200 mM H<sub>2</sub>O<sub>2</sub> (H) stained with nitroblue tetrazolium. The strength of the color showed the concentration of H<sub>2</sub>O<sub>2</sub> in the root tips.

(I) Total root length of JN177 and SR3 under stress conditions.

(J) The root H<sub>2</sub>O<sub>2</sub> content in the above seedlings. All data are given as mean ± sd. The double asterisks represent significant differences as determined by Student's *t* test at *P* < 0.01 (*n* = 60). FW, fresh weight.

Bars = 2 cm in (A) to (D) and 1 mm in (E) to (H).

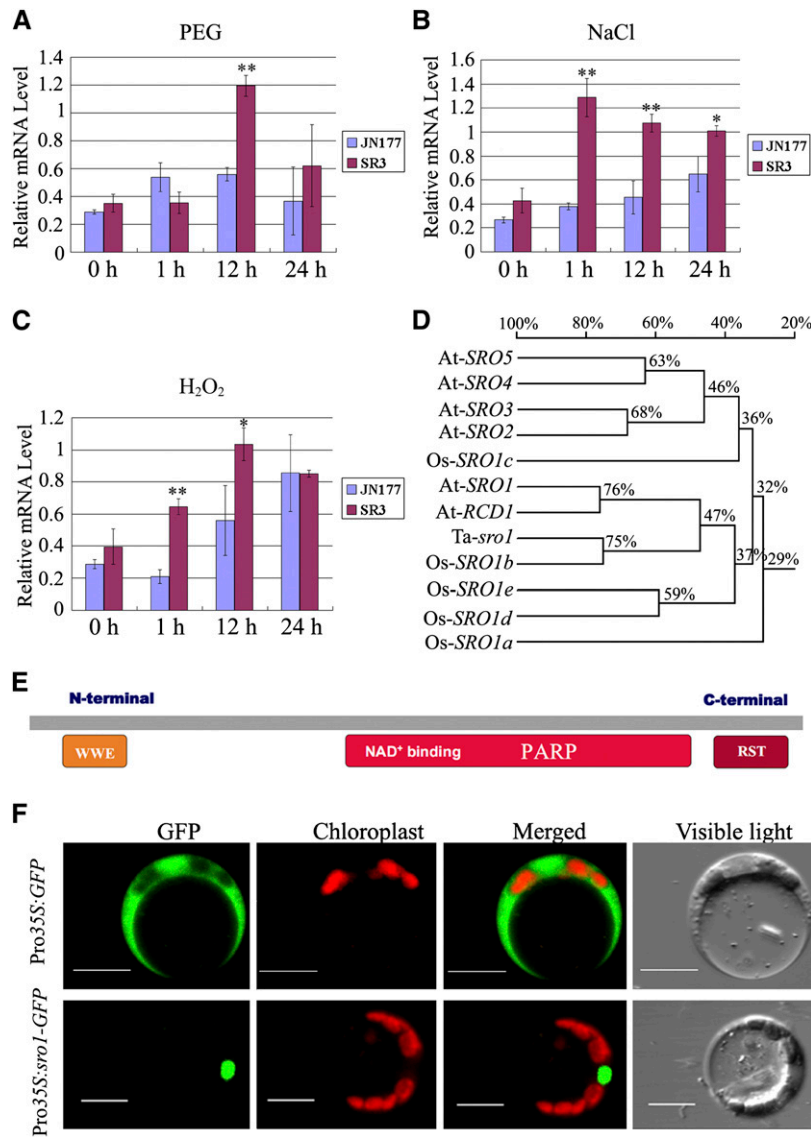
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of stress (18% PEG, 200 mM NaCl, and 200 mM H<sub>2</sub>O<sub>2</sub> applied for SR3 and JN177), a lower level (10% PEG, 100 mM NaCl, and 150 mM H<sub>2</sub>O<sub>2</sub>) was applied to test these lines. The constitutive expression of *Ta-sro1* increased the plant height as well as the root length under both control and stress conditions (Figures 3A to 3D and 3I; Supplemental Figure 4A), while its abolition had the opposite effect (Figures 3E to 3H and 3J; Supplemental Figure 4B). Accordingly, the ROS content of the former set of plants was higher (Figures 3K to 3N and 3S; Supplemental Figure 4C) and that of the latter was lower than the nonstressed controls (Figures 3O to 3R and 3T; Supplemental Figure 4D). Our investigations indicate that *Ta-sro1* was largely responsible for the seedling's vigorous growth and the stress tolerance of SR3.

### The Ta-sro1 Protein Has PARP Activity

Unlike well-studied PARP enzymes, such as *Homo sapiens* Hs-PARP1, Hs-PARP2, At-PARP1, and At-PARP2, which showed conserved PARP activities, some PARP catalytic domain-containing

proteins such as At-RCD1 did not possess the PARP activity (Jaspers et al., 2010). To further characterize the function mechanisms of the PARP catalytic domains of the SR3 *Ta-sro1* and JN177 *Ta-SRO1* proteins, we constructed three-dimensional models of SR3 *Ta-sro1*, JN177 *Ta-SRO1*, At-RCD1, At-PARP1, and At-PARP2 and compared them together with the reported crystal structure of Hs-PARP1 (PDB code: 1UK0) (Kinoshita et al., 2004). The resulting models were evaluated by two methods to make an integrative assessment of the structure quality, considering the geometries, stereochemistries, and energy distributions. The templates used for modeling these catalytic domains are shown in Supplemental Table 2. The evaluation results are indicative of a good quality of all five models (Supplemental Table 2). A structural analysis of Hs-PARP1 has shown that the active site pocket of its catalytic domain is negatively charged (Figure 4A) and includes the specific catalytic triad His<sup>862</sup>-Tyr<sup>896</sup>-Glu<sup>988</sup>, which is conserved in all PARP enzymes (Citarelli et al., 2010). These homology models revealed that the catalytic domains of *Ta-SRO1*, *Ta-sro1*, At-PARP1, and At-PARP2 possess an analogous negatively charged pocket



**Figure 2.** Expression, Gene Structure, and Subcellular Localization of *Ta-sro1*.

(A) to (C) Analysis of transcript differences between SR3 and JN177 exposed to 18% PEG (A), 200 mM NaCl (B), or 200 mM H<sub>2</sub>O<sub>2</sub> (C) by quantitative RT-PCR. All data are given as mean  $\pm$  SD of three independent biological replicates. The asterisks and double asterisks represent significant differences as determined by Student's *t* test at  $P < 0.05$  and  $P < 0.01$ , respectively.

(D) Phylogenetic analysis of *Ta-sro1* and its homologous genes. This tree is based on a multiple alignment that includes the full-length cDNA sequences. The percentage given on the phylogeny shows the similarity between sequences. Alignments used to generate the phylogeny are shown in Supplemental Data Set 1.

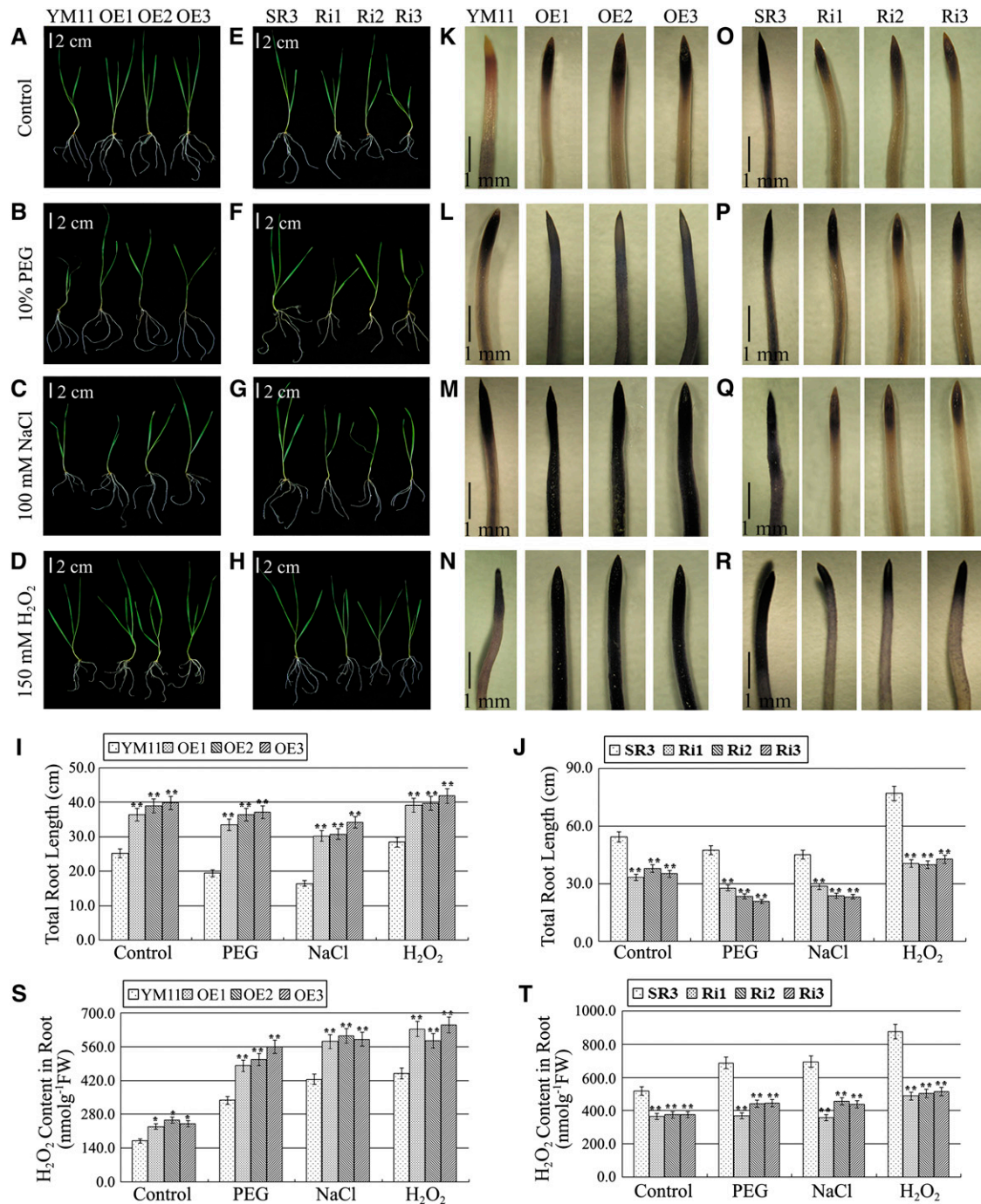
(E) Predicted structure of *Ta-sro1*. *Ta-sro1* contains the PARP signature and two protein-protein interaction domains: WWE and RST.

(F) Subcellular localization of *Ta-sro1* protein. Images were captured using the following wavelengths: GFP (excitation, 488 nm; emission, 509 nm) and chlorophyll autofluorescence (excitation, 448 nm; emission, 647 nm). Bars = 10  $\mu$ m.

(Figures 4B to 4E) to hold the positive NAD<sup>+</sup> substrate, carrying the catalytic triads Leu<sup>312</sup>-His<sup>344</sup>-His<sup>407</sup>, Leu<sup>312</sup>-His<sup>344</sup>-His<sup>407</sup>, His<sup>833</sup>-Tyr<sup>867</sup>-Glu<sup>960</sup>, and His<sup>486</sup>-Tyr<sup>520</sup>-Glu<sup>614</sup>, respectively. The corresponding pocket of At-RCD1 is positively charged and contains the unusual triad variant Leu<sup>345</sup>-His<sup>377</sup>-Asn<sup>440</sup> (Figure 4F), which may well explain its lack of catalytic ability. In addition, the predicted structure of *Ta-sro1* includes an arc-shaped face on the

opposite side of the catalytic domain, allowing for the localized accumulation of positive charge. The shape, size, and charge distribution over this surface of the predicted structure of *Ta-sro1* imply that it could function as a DNA binding site (Figure 4G).

To determine whether *Ta-SRO1* and *Ta-sro1* have PARP enzyme activity in vitro, we expressed and purified the *Ta-SRO1*-His<sub>6</sub>, *Ta-sro1*-His<sub>6</sub>, *Ta-sro1* (V250G)-His<sub>6</sub>, and *Ta-sro1*



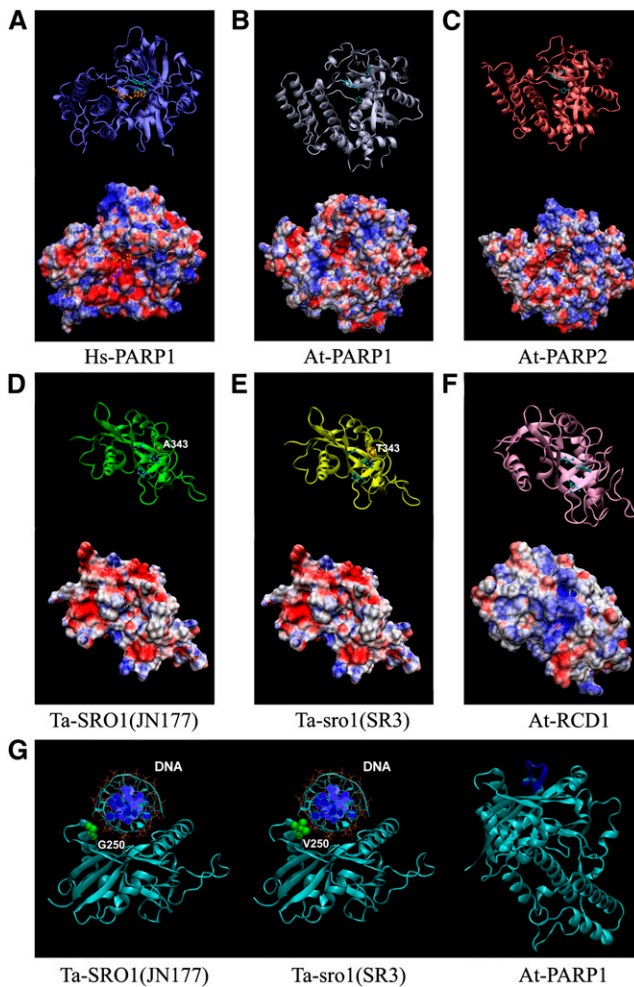
**Figure 3.** The Phenotype of Transgenic Wheat Lines under Abiotic Stress Treatment at the Seedling Stage.

(A) to (D) Phenotype of 1-week-old transgenic plants compared with YM11 under control (A), 10% PEG (B), 100 mM NaCl (C), or 150 mM H<sub>2</sub>O<sub>2</sub> (D). OE1, OE2, and OE3 are transgenic wheat lines constitutively expressing *Ta-sro1* cDNA from SR3 in wheat cultivar YM11.

(E) to (H) Phenotype of 1-week-old transgenic plants compared with SR3 under control (E), 10% PEG (F), 100 mM NaCl (G), or 150 mM H<sub>2</sub>O<sub>2</sub> (H). Nonstressed or stressed seedlings grown for 2 weeks. Ri1, Ri2, and Ri3 are independent RNAi transgenic wheat lines knocked down for the expression of *Ta-sro1* cDNA from SR3 in wheat cultivar SR3. YM11 and SR3 are nontransgenic controls.

(I) and (J) Total root length of the above seedlings.

(K) to (N) Comparison of root tips of 1-week-old seedlings under control (K), 10% PEG (L), 100 mM NaCl (M), or 150 mM H<sub>2</sub>O<sub>2</sub> (N) stained with nitroblue tetrazolium between *Ta-sro1* OE transgenic plants and YM11.



**Figure 4.** Structural Models of the Catalytic Domains of PARP-Containing Proteins.

(A) to (F) Top diagrams: Representations of the protein backbone of Hs-PARP1 (A), At-PARP1 (B), At-PARP2 (C), Ta-SRO1 (JN177) (D), Ta-sro1 (SR3) (E), and At-RCD1 (F) with the catalytic triad residues labeled with light-blue spheres and sticks. The inhibitor molecule of human Hs-PARP1 (A) is shown by orange spheres and sticks. The Ala-343 residue of Ta-SRO1 (D) and Thr-343 residue of Ta-sro1 (E) are marked by red spheres and sticks. Bottom diagrams: molecular surface electrostatic potential representations. Positively charged areas shown in blue and negatively charged ones in red.

(G) Hypothetical models of interactions between the catalytic domain and DNA. The Gly-250 of Ta-SRO1 and Val-250 of Ta-sro1 on the lateral protrusion of the arc-shaped face are marked by green spheres.

(T343A)-His<sub>6</sub> recombinant proteins from *Escherichia coli* cells (Figures 5A and 5B). The affinity-purified Ta-sro1 protein showed a higher level of PARP activity than did Ta-SRO1 (Figure 5C). However, reduced PARP activity could be detected in Ta-sro1 (V250G) and Ta-sro1 (T343A) (Figure 5C), in which the Val residue (V) at position 250 and Thr residue (T) at the position 343 of SR3 Ta-sro1 were replaced with the JN177 Ta-SRO1 residue Gly (G) and Ala (A), respectively. To demonstrate the ability of recombinant Ta-sro1-His<sub>6</sub> and Ta-SRO1-His<sub>6</sub> protein to bind to the damaged DNA in vitro, a dot blotting assay using a biotin-streptavidin system was performed. As shown in Figure 5D, Ta-sro1 exhibited significant binding toward the calf thymus DNA fragments. By contrast, Ta-SRO1 only showed weak binding to DNA fragments (Supplemental Figure 5). Taken together, these data indicate that SR3 Ta-sro1 is indeed a functional PARP enzyme and it shows higher activity than JN177 Ta-SRO1 due to the two point mutations.

To investigate whether the PARP activity of Ta-sro1 is required for its biological functions in vivo, we systematically compared the PARP activity between different plant combinations. PARP activity in SR3 was considerably higher than that in JN177 (Figure 5E), a result mirrored by the behavior of the transgenic Yangmai 11 constitutively expressing Ta-sro1 and the SR3 Ta-sro1 RNAi knockdown lines (Figures 5F and 5G). These results suggest that both overall growth vigor and stress tolerance may result from the PARP activity of SR3 Ta-sro1.

#### Constitutive Expression of Ta-sro1 Increased ROS Production and Enhanced Stress Tolerance in *Arabidopsis*

To gain further insight into the function of Ta-sro1 and the possible signaling pathways in which it participates, constitutive lines of Ta-sro1 were generated in *Arabidopsis* (Ta-sro1 overexpression [OE] lines). Two lines that accumulated detectable amounts of Ta-sro1 message (Supplemental Figure 6) were selected for further analysis. Root lengths of 1-week-old, 3-week-old and 8-week-old Ta-sro1 OE seedlings were substantially enhanced compared with those in the transgenic lines carrying an empty vector control (VC) lines (Figures 6A to 6C), which is caused by an enhanced cell division rate and larger cell size, as determined by comparing the cell numbers in the elongation zone and the root cell length (Figures 6D to 6F). The Ta-sro1 OE lines also displayed a notable improvement in their stress tolerance. This was particularly demonstrated by their superior root growth in the presence of 300 mM mannitol, 100 mM NaCl, 1.5 mM H<sub>2</sub>O<sub>2</sub>, and 1 μM MV, respectively (Figures 6G to 6K). A colorimetric assay showed that the Ta-sro1 OE lines accumulated more ROS than the VC lines under both nonstressed and stressed conditions (Figure 6L). Additionally, the Ta-sro1 OE lines exhibited more PARP activity

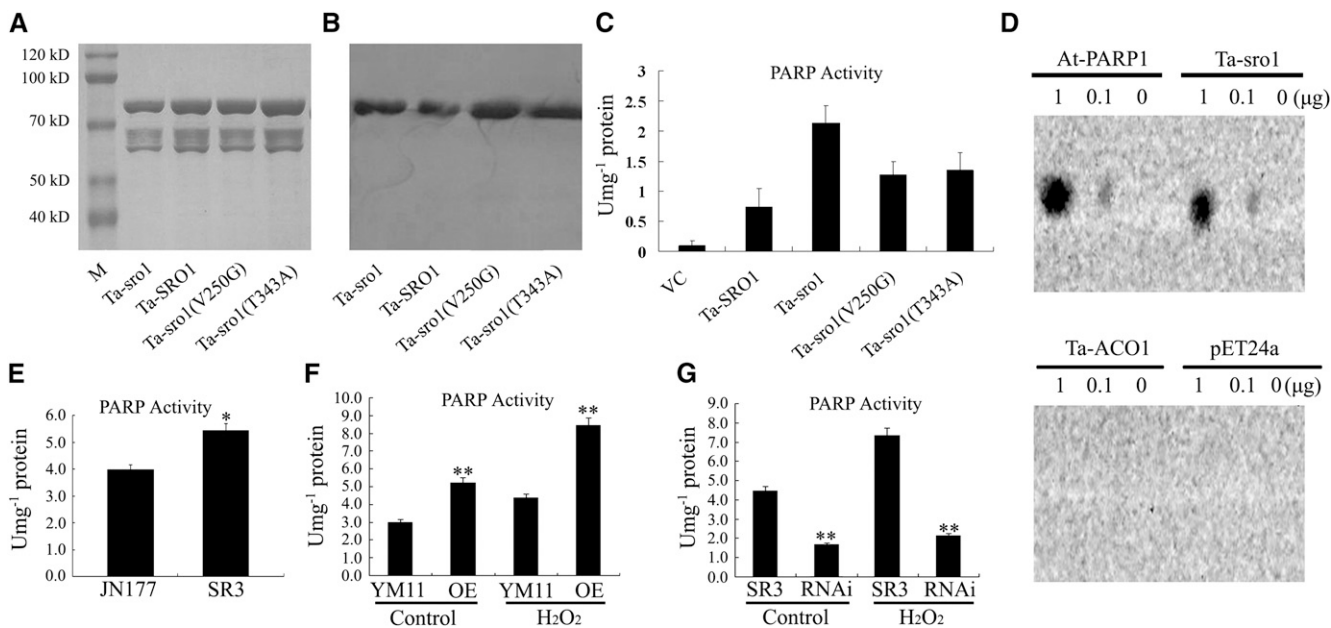
**Figure 3.** (continued).

(O) to (R) Comparison of root tips of 1-week-old seedlings under control (O), 10% PEG (P), 100 mM NaCl (Q), or 150 mM H<sub>2</sub>O<sub>2</sub> (R) stained with nitroblue tetrazolium between Ta-sro1 RNAi transgenic plants and SR3. The strength of the color showed the concentration of H<sub>2</sub>O<sub>2</sub> in the root tips.

(S) and (T) The root H<sub>2</sub>O<sub>2</sub> content in the above seedlings. All data are given as mean ± sd. The double asterisks represent significant differences determined by Student's *t* test at *P* < 0.01 (*n* = 15). FW, fresh weight.

Bars = 2 cm in (A) to (H) and 1 mm in (K) to (R).

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**Figure 5.** The in Vitro and in Vivo Poly(ADP Ribosyl)ation Activity of Ta-sro1.

**(A)** Photograph of purified Ta-SRO1-His<sub>6</sub>, Ta-sro1-His<sub>6</sub>, Ta-sro1 (V250G)-His<sub>6</sub>, and Ta-sro1 (T434A)-His<sub>6</sub> separated by SDS-PAGE and stained with Coomassie Brilliant Blue R 250. M, Blue Plus II protein marker (TransGen Biotech).

**(B)** Immunoblot analysis of the above purified proteins. Different His fusion proteins were detected using an anti-His antibody.

**(C)** In vitro PARP activity assay. Total proteins from *E. coli* cells containing pET24a VC induced for 3 h by 1 mM isopropyl β-D-1-thiogalactopyranoside were analyzed for comparison.

**(D)** In vitro DNA binding activity of Ta-sro1-His<sub>6</sub>. Purified At-PARP1 protein was used as the positive control, while purified Ta-ACO1-His<sub>6</sub> (for wheat aminocyclopropane-1-carboxylate oxidase 1) protein and pET24a total protein were used as negative controls.

**(E)** PARP activity in JN177 and SR3.

**(F)** PARP activity in wheat plants constitutively expressing Ta-sro1. The PARP activity of Ta-sro1 OE lines was higher than the nontransgenic control YM11 both under stress (YM11-H<sub>2</sub>O<sub>2</sub> and OE-H<sub>2</sub>O<sub>2</sub>) and nonstress (YM11-C and OE-C) conditions.

**(G)** PARP activity in wheat RNAi knockdown plants of Ta-sro1. The PARP activity of Ta-sro1 RNAi lines was lower than that of SR3 both under stress (SR3-H<sub>2</sub>O<sub>2</sub> and OE-H<sub>2</sub>O<sub>2</sub>) and nonstress (SR3-C and RNAi-C) conditions. All data are given as mean ± SD. The asterisks and double asterisks represent significant differences determined by Student's *t* test at *P* < 0.05 and *P* < 0.01, respectively.

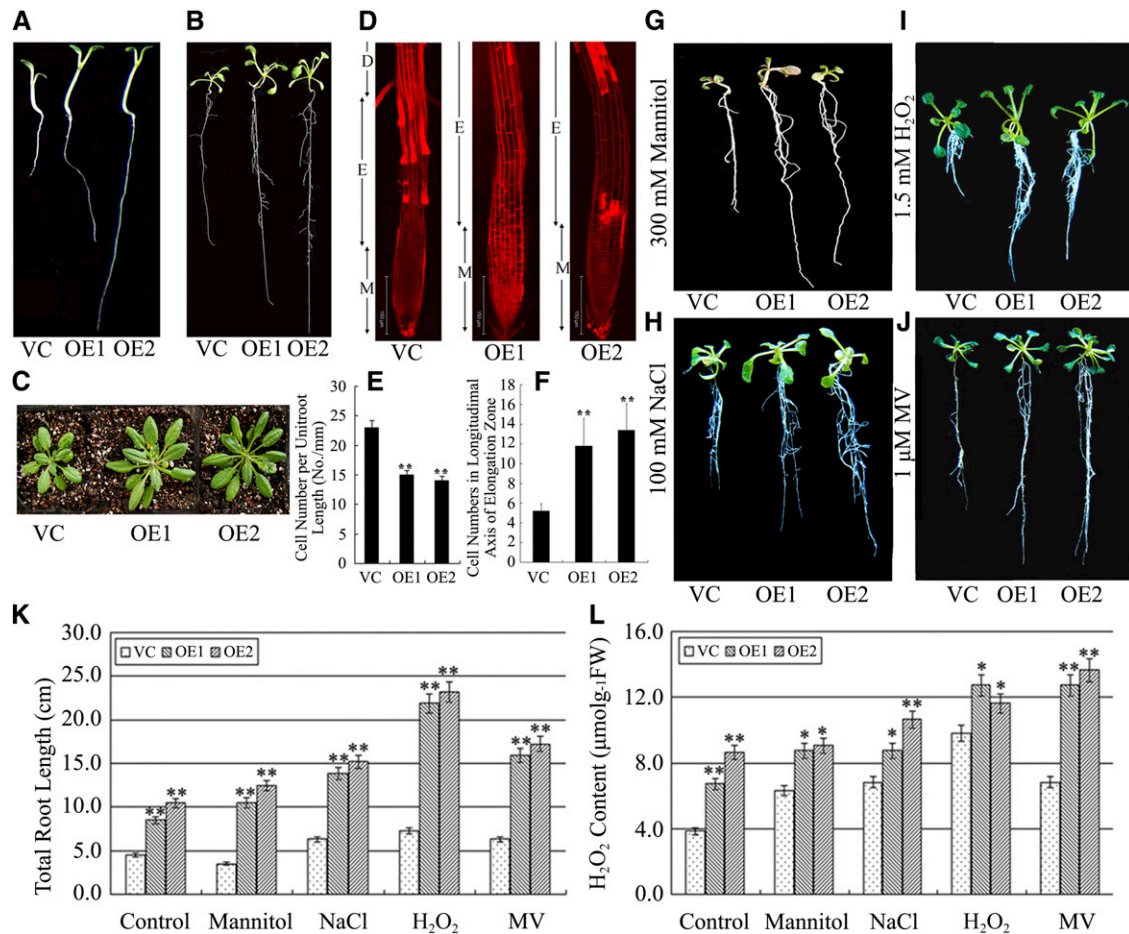
than did the VC plants (Figure 7A), and root elongation was more severely compromised by the presence of the PARP inhibitor nicotinamide (NA) (Figures 7B to 7D), which further proved that these phenotypes conferred by the Ta-sro1 variant are not simply due to its higher level of expression but due to its higher PARP activity. Collectively, these data indicated that the heterologous expression of Ta-sro1 in *Arabidopsis* showed consistent phenotypes with wheat Ta-sro1 OE plants and SR3 and thus that *Arabidopsis* Ta-sro1 OE lines can be used for further research into the mechanism by which Ta-sro1 enhances seedling growth and abiotic stress resistance.

To confirm that Ta-sro1 enhances stress tolerance by increasing ROS level, we introduced the Pro35S:sro1 transgene into the *At-rbohF* mutant background, in which the primary root growth is repressed (Kwak et al., 2003). We found that Ta-sro1 could rescue the shorter root of *At-rbohF* (Supplemental Figure 7) under nonstressful conditions. When exposed to 0.6 mM H<sub>2</sub>O<sub>2</sub>, the root elongation of the *At-rbohF* mutant was more severely inhibited, whereas Ta-sro1 overexpression in the *At-rbohF*/35S:sro1 transgenic line rescued the root length of the *At-rbohF* mutant, resulting in plants with roots that were even a little longer than those

of the wild type (Supplemental Figure 7). Consistently, the Ta-sro1 OE line showed the longest root (Supplemental Figure 7). These results indicated that the higher ROS levels of Ta-sro1 OE lines was responsible for its stress tolerance.

#### ROS Accumulation in the *Arabidopsis* Ta-sro1 OE Lines

As NADPH oxidase (NOX) activity has been implicated in ROS formation (Foreman et al., 2003; Mittler et al., 2004) and Ta-sro1 can complement the *At-rbohF* (an important gene of NOX) mutant phenotype, NOX activity was compared between SR3 and JN177. Consistent with their contrasting accumulation of ROS, NOX activity was higher in the former (Figure 8A). Consistent with our expectation, the Ta-sro1 OE lines also showed higher NOX activity than VC lines (Figure 8B). When the plants were treated with the NOX inhibitor diphenylene iodonium (DPI), a measurable decrease in ROS accumulation and root elongation was observed for the Ta-sro1 OE lines (Figures 8C to 8H). A further source of ROS (particularly H<sub>2</sub>O<sub>2</sub>) is the mitochondrial electron transport chain, which in plants includes alternative oxidase (AOX) and various NAD (P)H dehydrogenases. The most ubiquitously expressed AOX gene



**Figure 6.** Phenotype of *Arabidopsis* Plants Constitutively Expressing *Ta-sro1*.

(A) to (C) Phenotypes of 1-week-old (A), 3-week-old (B), and 8-week-old (C) transgenic plants compared with VC plants. The VC plants are a transgenic line carrying an empty vector. OE1 and OE2 are transgenic *Arabidopsis* lines constitutively expressing *Ta-sro1* cDNA from SR3. (D) Root tips of 1-week-old seedlings as visualized by confocal laser scanning microscopy after staining with propidium iodide. The elongation zone of OE lines was significantly longer than that of the VC. M, meristematic zone; E, elongation zone; D, differentiation zone. (E) Cell number statistics in roots of 1-week-old seedlings, where the larger size of OE line root elongation zone cells resulted in fewer cells per unit root length. (F) Cell number statistics in the longitudinal axis of the root elongation zone of 1-week-old seedlings. The elongation zone of OE lines contains more cells than that of the VC. (G) to (J) Phenotype of 1-week-old transgenic seedlings stressed for 2 weeks with either 300 mM mannitol (G), 100 mM NaCl (H), 1.5 mM H<sub>2</sub>O<sub>2</sub> (I), or 1 μM MV (J) compared with the VC. (K) and (L) Total root length (K) and root H<sub>2</sub>O<sub>2</sub> content (L) in stressed seedlings. All data are given as mean ± sd. The asterisks and double asterisks represent significant differences, as determined by Student's *t* test at  $P < 0.05$  and  $P < 0.01$ , respectively ( $n = 30$ ). FW, fresh weight.

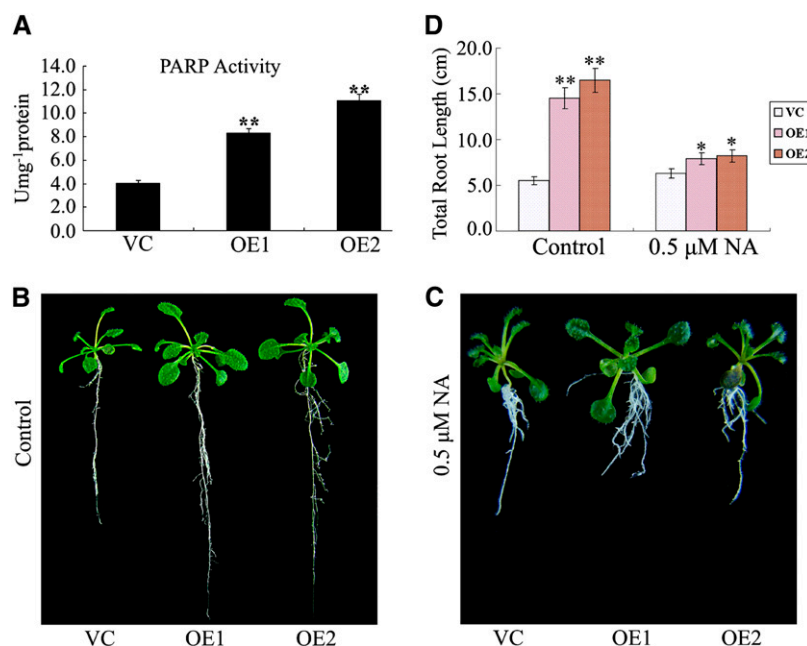
in *Arabidopsis* is *At-AOX1a*, while *At-NDB2* encodes an NAD(P)H dehydrogenase frequently coexpressed with *At-AOX1a* as part of the stress response (Clifton et al., 2006). Expression analysis indicated that *At-AOX1a* was more markedly downregulated in the *Ta-sro1* OE line than in the VC plants, while *At-NDB2* was upregulated (Figures 8I and 8J). The AOX-dependent respiration rate was measured either in the presence of potassium cyanide (KCN; a cytochrome inhibitor) or salicyl hydroxamic acid (SHAM; an AOX inhibitor). No significant difference between respiration rate of *Ta-sro1* OE and VC was detectable prior to inhibitor treatment, but respiration was significantly increased in the former in the presence of SHAM and reduced by KCN (Figure 8K), suggesting

that the *Ta-sro1* transgene downregulates AOX. Furthermore, the LD<sub>50</sub> (half lethal dose of KCN) for KCN was 150 μM for the *Ta-sro1* OE lines but 300 μM for VC lines (Figure 8L). Taken together, the enhanced NOX activity and altered expression of *At-AOX1a* and *At-NDB2* genes in the *Ta-sro1* OE lines may be responsible for higher ROS accumulation in roots.

#### ROS Scavenging in the *Arabidopsis Ta-sro1* OE Lines

To address the possibility that ROS were detoxified more rapidly in the *Ta-sro1* OE lines or that metabolism of these lines is more tolerant of elevated ROS content, the activities of various key





**Figure 7.** PARP Activity and the Effect of the PARP Inhibitor NA on Root Growth in *Arabidopsis* Ta-*sro1* OE Lines.

(A) Comparison of PARP activity between VC and Ta-*sro1* OE lines.

(B) and (C) Seedling phenotype of transgenic *Arabidopsis* lines compared with control plants under normal conditions (B) and 0.5 μM NA treatment (C).

(D) Effect of NA treatment on root growth. All data are given as mean ± SD. The asterisks and double asterisks represent significant differences, as determined by Student's *t* test at  $P < 0.05$  and  $P < 0.01$ , respectively ( $n = 30$ ).

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enzymes (superoxide dismutase [SOD], peroxidase [POD], catalase [CAT], ascorbate peroxidase [APX], and glutathione peroxidase [GPX]) were determined. No significant difference was noted between the Ta-*sro1* OE and VC lines with respect to either SOD or POD activity, but the level of both APX and GPX (involved in the removal of hydrogen peroxide) activity was higher in the Ta-*sro1* OE lines, and CAT activity was lower (Figures 9A to 9C). To characterize ROS scavenging enzymes in SR3 and JN177, we also determined the activity of some key enzymes in wheat. Consistent with the enzyme activities in the Ta-*sro1* OE and VC lines, SR3 also showed higher APX and lower CAT activity than JN177 (Figures 9D and 9E). No significant difference in SOD and POD activity was found between JN177 and SR3. These results suggest that the Ta-*sro1* OE lines were more effective than the VC lines in terms of ROS detoxification.

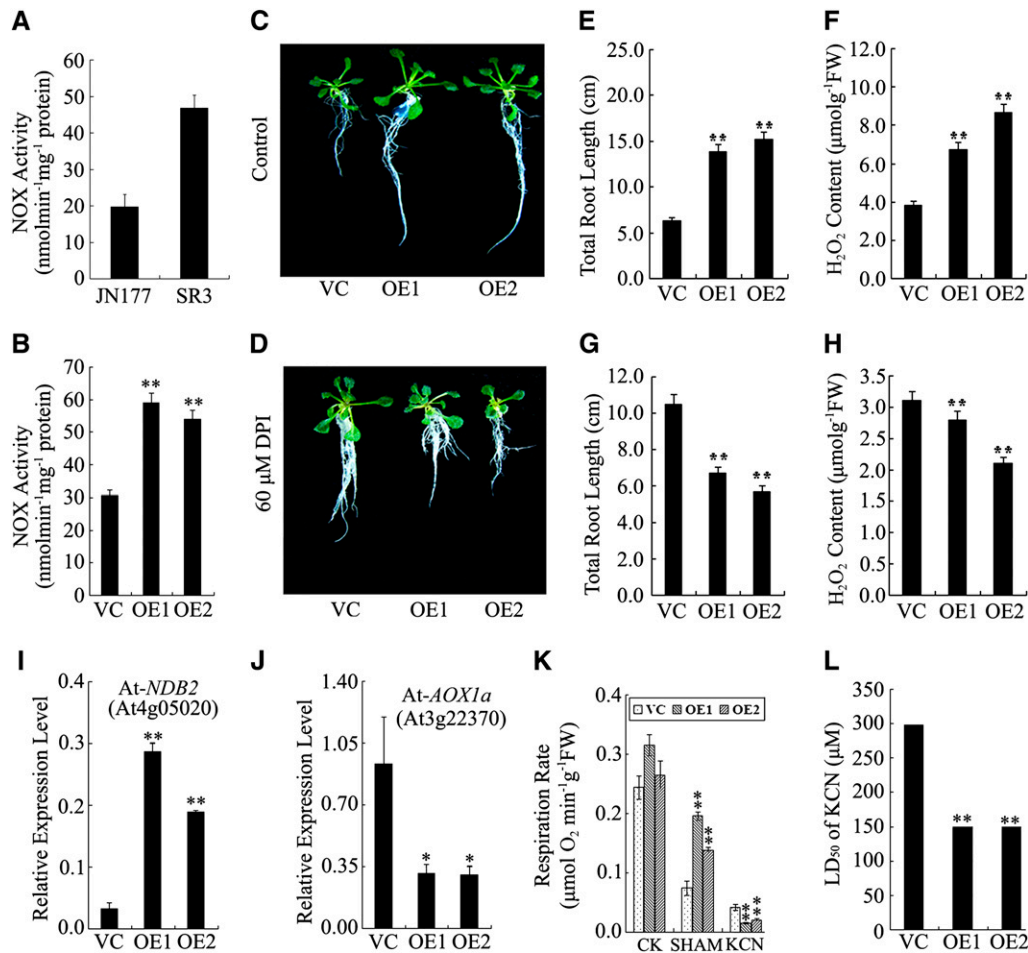
The same approach was taken regarding redox balance maintenance, where the activities of enzymes in the ascorbate (AsA)-GSH cycle (monodehydroascorbate reductase [MDAR], dehydroascorbate reductase [DHAR], and glutathione reductase [GR]) and the presence of the four antioxidant couples AsA/dehydroascorbate (DHA), GSH/GSSG, NADH/NAD<sup>+</sup>, and NADPH/NADP<sup>+</sup> were monitored. MDAR, DHAR, and GR activity was greater in the Ta-*sro1* OE lines than in the VC lines (Figures 9F to 9H). While the AsA/DHA ratio was maintained more effectively in the Ta-*sro1* OE lines, those for NADH/NAD<sup>+</sup>, NADPH/NADP<sup>+</sup>, and GSH/GSSG ratios were all reduced (Figures 9I to 9L). Thus, we can conclude that Ta-*sro1* promotes the activity of AsA-GSH cycle enzymes and GPX cycle enzymes, which control ROS content and cellular redox homeostasis.

### Increased Genomic Integrity in the *Arabidopsis* Ta-*sro1* OE Lines

PARPs have been implicated in DNA damage repair and genomic integrity (Kim et al., 2005; Schreiber et al., 2006; Hassa and Hottiger, 2008). The involvement of Ta-*sro1* in ROS homeostasis, PARP activity, and the nuclear localization of Ta-*sro1* prompted us to test whether Ta-*sro1* also functions in this process. We therefore detected the damage to genomic DNA of Ta-*sro1* OE and VC lines under UV irradiation and H<sub>2</sub>O<sub>2</sub> treatments by single-cell gel electrophoresis (SCGE). As shown in Figure 10, Ta-*sro1* OE protoplasts were more resilient than those of the VC. Similarly, when exposed to UV irradiation, the protoplasts of SR3 showed less DNA damage than JN177 (Supplemental Figure 8), which further proved that Ta-*sro1* may be largely responsible for the stress tolerance of SR3. Taken together, these results imply that Ta-*sro1* plays an important role in DNA damage repair and genomic integrity through its PARP activity, by which it copes with the abiotic stresses and maintains plant growth.

### DISCUSSION

The PARP proteins are widely distributed among the eukaryotes (although are absent from yeast) and appear to be involved in processes ranging from DNA repair and the regulation of chromatin and telomere structure to transcriptional regulation, the response to abiotic and biotic stress, and the activation of cell death (Beneke and Bürkle, 2007; Citarelli et al., 2010). Surprisingly, this important



**Figure 8.** Characterization of ROS Production Elements in Wheat and *Arabidopsis* Plants Constitutively Expressing *Ta-sro1*.

(A) Comparison of NOX activity between JN177 and SR3.

(B) Comparison of NOX activity between *Ta-sro1* OE lines and VC plants.

(C) and (D) Phenotype of VC, *Ta-sro1* OE1, and *Ta-sro1* OE2 transgenic plants grown in normal conditions (C) or in the presence of 60 μM NOX inhibitor DPI (D).

(E) to (H) Root growth and H<sub>2</sub>O<sub>2</sub> content in seedlings grown in the absence (E) and (F) or presence of DPI (G) and (H). FW, fresh weight.

(I) and (J) Expression levels of *At-NDB2* [NAD(P)H dehydrogenase] (I) and *At-AOX1a* (J) between *Ta-sro1* OE lines and VC plants.

(K) and (L) respiration rate (K) and LD<sub>50</sub> (half lethal dose of KCN) for KCN (L) between *Ta-sro1* OE lines and VC plants. All data are given as mean ± SD. The double asterisks represent significant differences, as determined by Student's *t* test at *P* < 0.01 (*n* = 30).

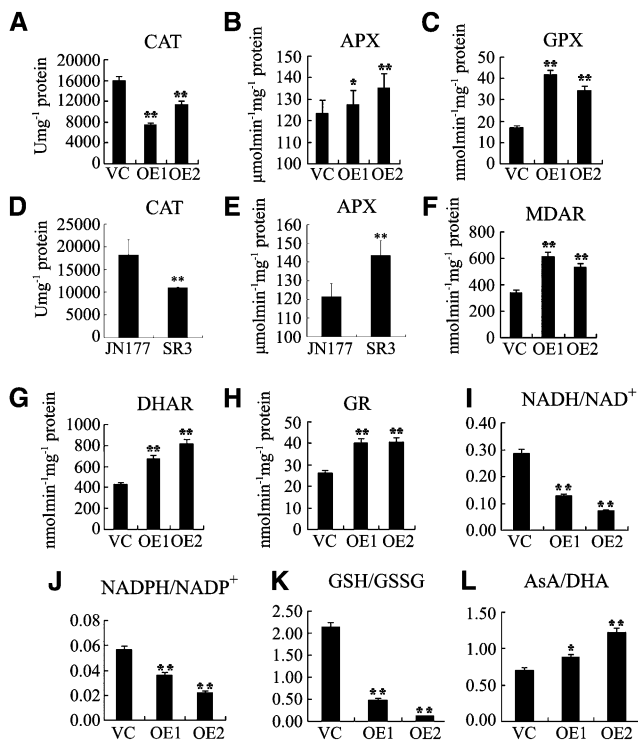
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protein family has received little attention from plant scientists to date (Skirycz et al., 2011). Here, we described the functional characterization of a wheat PARP and showed that it underpins both seedling vigor and tolerance to abiotic stress, through its catalytic activity and modulation of redox homeostasis.

### The PARP Activity of *Ta-sro1* in Relation to the DNA Integrity of *Ta-sro1* OE Lines

Eukaryotic organisms express multiple PARP proteins, which although recognized by the presence of a conserved PARP catalytic domain, do not all display PARP activity (Citarelli et al., 2010). The presence of a negatively charged pocket able to retain the NAD<sup>+</sup> substrate, together with its LHH catalytic triads (although not a

classical one) (Figure 4E), suggested that SR3 *Ta-sro1* would have PARP activity, which was borne out experimentally (Figure 5C). The greater PARP activity of *Ta-sro1* may well flow from substitution from Ala-343 to Thr-343 of *Ta-sro1* because the hydrophobic Ala residue does not contribute to the interaction with the substrate, whereas the hydrophilic Thr residue does so by presenting both a methyl and a hydroxyl group to the catalytic surface, which encourages interaction with the substrate (Figures 4D and 4E). Unlike the well-characterized animal and plant PARPs, which feature at least one DNA binding domain, SRO proteins have no such domains (Citarelli et al., 2010). The predicted structure of *Ta-sro1* includes an arc-shaped face on the far side of the catalytic domain (opposite to the catalytic pocket), allowing for the localized accumulation of positive charge. The shape, size,



**Figure 9.** Cellular Redox Balance Is Affected in Wheat and *Arabidopsis* Plants Constitutively Expressing *Ta-sro1*.

(A) to (C) Comparison of enzyme activities of CAT (A), APX (B), and GPX (C) between *Ta-sro1* OE lines and VC plants.

(D) and (E) Comparison of enzyme activities of CAT (D) and APX (E) between JN177 and SR3.

(F) to (H) Comparison of enzyme activities of MDAR (F), DHAR (G), and GR (H) between *Ta-sro1* OE lines and VC plants.

(I) to (L) Comparison of the ratios of NADH/NAD<sup>+</sup> (I), NADPH/NADP<sup>+</sup> (J), GSH/GSSG (K), and AsA/DHA (L) between *Ta-sro1* OE lines and VC plants. All data are given as mean  $\pm$  sd. The asterisks and double asterisks represent significant differences, as determined by Student's *t* test at  $P < 0.05$  and  $P < 0.01$ , respectively.

and charge distribution over this surface of the predicted structure of *Ta-sro1* suggested that this surface may function as a DNA binding site (Figure 4G); therefore, it is reasonable to find that the *Ta-sro1* protein has DNA binding ability (Figure 5D). Intriguingly, a second allelic difference between the SR3 *Ta-sro1* and JN177 *Ta-SRO1* proteins (Val-250/Gly-250) lies on a lateral protrusion from this surface (Figure 4G), which implies that the more hydrophobic Val in the SR3 version of the protein increases its DNA binding capacity over that achieved by the JN177 version (Supplemental Figure 5). This may contribute to the fact that *Ta-sro1* (V250G) protein has decreased PARP enzyme activity compared with *Ta-sro1* (Figure 5C). By contrast, Hs-PARP and At-PARP both comprise a two-stranded  $\beta$ -sheet above the arc-shaped face, giving rise to a structure that mitigates against DNA binding at this site (Figure 4G).

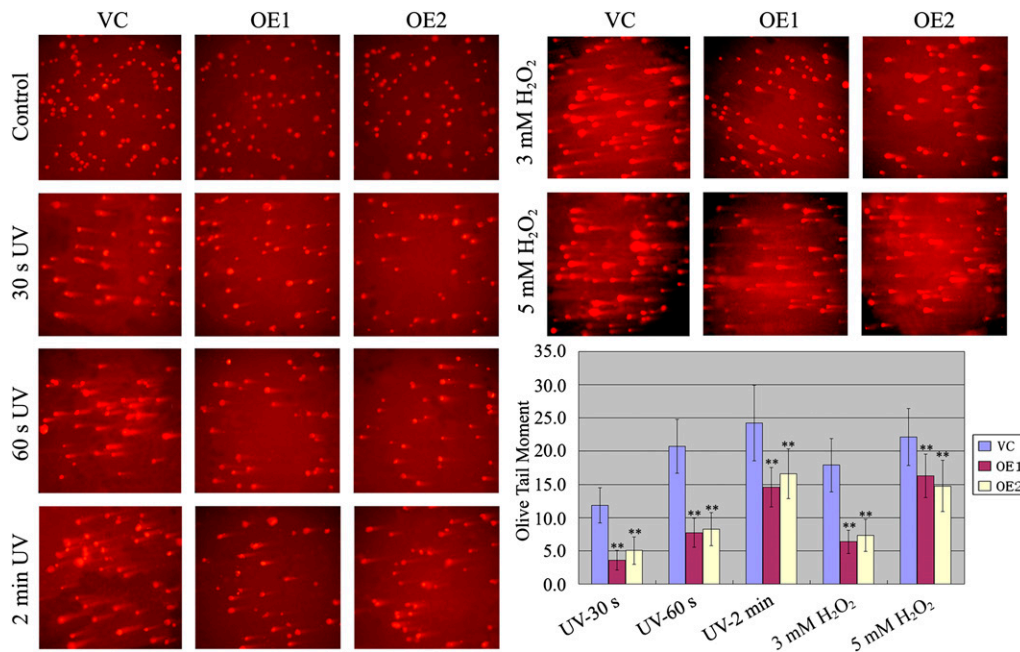
Reduced PARP activity has been reported to improve tolerance to a broad range of abiotic stresses, through its reduction in NAD<sup>+</sup> consumption (De Block et al., 2005; Vanderauwera et al., 2007). The strong activation of PARP and poly(ADP ribosylation) can alter

the cellular redox state, leading to a depletion in the availability of ATP, which in severe cases induces cell death (Schraufstatter et al., 1986). On the contrary, the constitutive expression of the SR3 allele of *Ta-sro1* in *Arabidopsis*, despite the increased PARP activity of its gene product (Figure 7A) and decreased ATP content (Supplemental Figure 9), enhanced both plant growth and stress tolerance. This is explainable. Under normal conditions, heterologous expression of SR3 *Ta-sro1* in *Arabidopsis* resulted in higher PARP activity and, thus, more ATP depletion. Upon exposure to abiotic stress (NaCl or H<sub>2</sub>O<sub>2</sub>), the ATP content of *Ta-sro1* OE lines increased compared with the decrease or less of an increase ATP level in the wild type (Supplemental Figure 9). It is therefore possible that *Ta-sro1* represses the alternative pathway by repressing *AOX1a* expression, thereby promoting the efficiency of ATP synthesis. This also implies that energy homeostasis is the basis of the stress tolerance.

PARP is also one of the earliest response factors to DNA breakage and plays an important role in maintaining DNA integrity (Kim et al., 2005). In mouse, both *Parp1* and *Parp2* knockouts are hypersensitive to alkylating agents as a result of their defective DNA excision repair machinery (Trucco et al., 1998; Schreiber et al., 2002; Kim et al., 2005; Schreiber et al., 2006; Hassa and Hottiger, 2008). The constitutive expression of *Arabidopsis Parp* in soybean (*Glycine max*) cells subjected to a mild level of oxidative stress reduced the frequency of ROS-induced DNA damage, while abolishing it produced the opposite effect (Amor et al., 1998). Here, SCGE analysis suggested that the higher PARP activity (Figure 7A) in *Ta-sro1* OE transgenic plants helped maintain DNA integrity by enhancing the activity of the DNA repair machinery (Figure 10). In mammalian cells, PARP-1 could interact directly with ATM (for Ataxia-Telangiectasia mutated), an important sensor of ROS (Guo et al., 2010), and form a large protein complex to function in DNA repair and maintenance of genomic integrity through its PARP activity (Aguilar-Quesada et al., 2007; Gagné et al., 2008). In our study, *At-ATM* had a higher expression level in *Arabidopsis Ta-sro1* OE lines than in the VC line under both NaCl and H<sub>2</sub>O<sub>2</sub> treatments (Supplemental Figure 10). In addition, the PARP domain of *Ta-sro1* is accompanied by two different putative protein-protein interaction domains, WWE and RST. Thus, *Ta-sro1* could possibly act as a scaffold protein that assembles DNA damage repair proteins for their posttranslational modification, relocalization, or degradation.

### ***Ta-sro1* Promotes Growth and Abiotic Stress Tolerance by Regulating Redox Homeostasis**

We demonstrated that the constitutive expression of *Ta-sro1* in wheat and *Arabidopsis* enhanced the accumulation of ROS and promoted plant growth and stress tolerance (Figures 3 and 6). The production of ROS by the NOX homolog (*Rboh*) was first discovered in the defense reactions against pathogens, and the discovery of new functions for plant *Rbohs* underlined diverse roles for NOX-generated ROS, including hormone synthesis, the regulation of stomatal closure, and root hair growth (Hammond-Kosack and Jones, 1996; Sagi and Fluhr, 2001; Foreman et al., 2003; Kwak et al., 2003). The discovery that NOX activity was substantially increased in SR3 and *Ta-sro1* OE lines (Figures 8A and 8B) and that *Ta-sro1* could complement the absence of *At-rbohF* (Supplemental Figure 7) suggested that NOX-generated ROS also



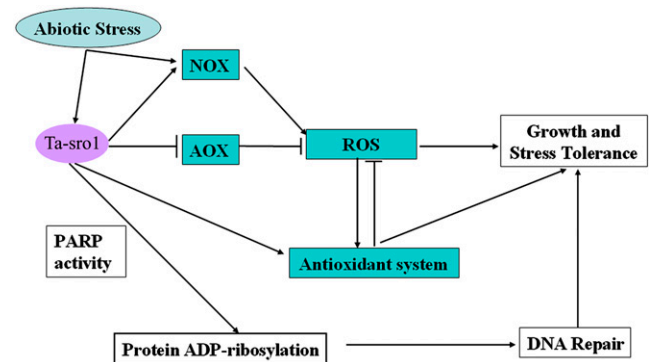
**Figure 10.** SCGE Analysis of the Protoplasts of Wild-Type and *Arabidopsis* Plants Constitutively Expressing *Ta-sro1* Exposed to UV or H<sub>2</sub>O<sub>2</sub>.

DNA damage in protoplasts of *Arabidopsis* plants constitutively expressing *Ta-sro1* and VC plants exposed to UV for 30 s, 60 s, and 2 min or 3 mM H<sub>2</sub>O<sub>2</sub> or 5 mM H<sub>2</sub>O<sub>2</sub> as indicated. Olive tail moment (OTM) represents the product of the percentage of total DNA in the tail (TD) and the tail length (TL): OTM = TL × TD. Abbreviations are as in Figure 5. All data are given as mean ± SD of four independent experiments. The double asterisks represent significant differences, as determined by Student's *t* test at *P* < 0.01.

acts in plant growth and abiotic stress. NAD(P)H dehydrogenase is a non-proton-pumping respiratory enzyme that functions in parallel with complex I to transfer electrons from NAD(P)H to ubiquinone. Together, the ubiquinone radical and NAD(P)H dehydrogenase are the main generators of ROS in the mitochondrion (Dat et al., 2000), so it is reasonable to suggest that in the *Ta-sro1* OE lines, the upregulation of the NAD(P)H dehydrogenase NDB2 (Figure 8H) would increase the production of ROS. In addition, the upregulation of both NOX and NAD(P)H dehydrogenase would likely decrease the NAD(P)H/NAD(P)<sup>+</sup> ratio via the oxidation of NAD(P)H (Figures 9I and 9J). AOX activity helps to regulate oxygen homeostasis within the mitochondria, and its suppression results in the elevation of ROS content (Maxwell et al., 1999). Both the gene expression and respiration inhibition experiments indicated a repression of AOX in the *Ta-sro1* OE lines (Figure 8), which can explain, at least in part, the extra accumulation of ROS.

The dual function of ROS implies that its cellular concentration has to be tightly controlled (Apel and Hirt, 2004). It has been proposed that AsA and GSH act as ROS scavengers either interdependently (AsA-GSH cycle) or independently (GPX cycle), and the enzymes include APX, GPX, MDAR, DHAR, and GR (Foyer and Noctor, 2011). It was possible to show that most of the extra ROS induced by the constitutive expression of *Ta-sro1* was scavenged through enzymes of the GPX cycle and AsA-GSH cycle and not by POD, SOD, or CAT (Figure 9). Note that both of these ROS processing pathways need GSH and NAD(P)H as reducing agents, thereby reducing the GSH/GSSG and NAD(P)H/NAD(P)<sup>+</sup> ratios (Figures 9I to 9K). The efficient antioxidant system ensures redox homeostasis, which can be indicated by the lower

malondialdehyde (MDA) content in *Ta-sro1* overexpression lines of *Arabidopsis* under salinity (Supplemental Figure 11), in agreement with that in SR3 (Liu et al., 2012).



**Figure 11.** A Proposed Model for the Role of *Ta-sro1* in the Regulation of Abiotic Stress Tolerance and Plant Growth.

Abiotic stress induced the expression of *Ta-sro1*, which promoted the accumulation of ROS, mainly by enhancing NOX activity and NAD(P)H dehydrogenase expression in conjunction with the suppression of AOX expression. At the same time, *Ta-sro1* promoted the activities of enzymes in the ROS scavenging system to regulate cellular redox homeostasis and maintain the ROS content to a higher level, through which stress tolerance and plant growth are controlled. Meanwhile, the PARP activity of *Ta-sro1* could enhance the genome stability of plants by modulating DNA damage repair, which also improves stress tolerance and plant growth.

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In conclusion, we demonstrated that Ta-sro1 is involved in seedling vigor and tolerance to abiotic stress, acting via the modulation of redox homeostasis and the maintenance of genome stability (Figure 11). A possible outline mechanism is that its constitutive expression enhances NOX activity and NAD(P)H dehydrogenase expression, while at the same time repressing AOX expression. As a result, the plant accumulates a higher level of H<sub>2</sub>O<sub>2</sub> than does the wild type. It also enhances the regulation exerted by the GPX cycle and AsA-GSH cycle over H<sub>2</sub>O<sub>2</sub> concentration and redox homeostasis. Meanwhile, the PARP enzyme activity of Ta-sro1 helps to maintain DNA integrity. Notably, based on the presented data we cannot exclude the possibility that Ta-sro1 acts by other mechanisms; however, all of the above properties make Ta-sro1 an excellent candidate for plant genetic improvement.

## METHODS

### Plant Materials and Growing Conditions

The bread wheat (*Triticum aestivum*) cultivar SR3 was derived from an asymmetric somatic hybrid between the cultivar JN177 and *Thinopyrum ponticum*. SR3, the salinity sensitive wheat cultivar Yangmai 11 and wild-type *Arabidopsis thaliana* ecotype Columbia-0 are all maintained in our laboratory. For germination, wheat grains were first surface sterilized in 2% (w/v) sodium hypochlorite (NaClO) and then laid on moistened filter paper for 2 d at 20°C. Seedlings of uniform size were transferred to a 12-h photoperiod regime (light/dark temperature 22/18°C) under 300  $\mu\text{mol m}^{-2} \text{s}^{-1}$  illumination and a relative humidity of 50% and were grown hydroponically in half-strength Hoagland solution. Abiotic stress was applied to SR3 and JN177 by the addition of 18% (w/v) PEG, 200 mM NaCl, or 200 mM H<sub>2</sub>O<sub>2</sub> to the hydroponic solution after 1 week. A lower level of abiotic stress was applied to the Ta-sro1 OE lines and RNAi lines together with control lines by the addition to the hydroponic solution of 10% (w/v) PEG, 100 mM NaCl, or 150 mM H<sub>2</sub>O<sub>2</sub> after 1 week. *Arabidopsis* seeds were surface sterilized by immersion in 0.1% (w/v) mercuric chloride and plated on a 1% agar medium supplemented with Murashige and Skoog salts and 3% (w/v) Suc. After keeping seeds in the dark at 4°C for 3 d, the plates were removed to a 16-h photoperiod, 22°C, and 70% relative humidity regime (light intensity 170 to 200  $\mu\text{mol m}^{-2} \text{s}^{-1}$ ). One-week-old seedlings were exposed to either 300 mM mannitol, 100 mM NaCl, 1.5 mM H<sub>2</sub>O<sub>2</sub>, 1  $\mu\text{M}$  MV, or 50 to 600  $\mu\text{M}$  KCN for 2 weeks.

### Transgenic Constructs of Ta-sro1 and Genetic Transformation

To generate the RNAi construct, we searched the wheat genome sequence database (<http://www.wheatgenome.org>) for SRO and PARP homologous genes and identified 10 wheat SRO and eight PARP homologous sequences. Among them, only three sequences (chr5bl\_10790093, chr4al\_7142517, and chr5dl\_4543936 referring to the three Ta-SRO1 alleles) located on 5B, 4A, and 5D genomes of wheat showed high similarity with Ta-sro1. The RNAi construct was designed in the 3' end of Ta-sro1, which contains the RST domain and 3'-untranslated region of the above 3 SRO1 sequences. A 410-bp fragment was amplified using primers Ta-sro1Ri-F and Ta-sro1Ri-R and inserted conversely into the vector pTCK303 (under the control of the *Zm-Ubiquitin* promoter) to knock down Ta-sro1 and the other two alleles of SR3.

To generate the Pro35S:sro1 construct for overexpression in *Arabidopsis* and the *At-rbohF* mutant, Ta-sro1 coding DNA sequence (CDS) was amplified with primers Ta-sro1AOE-F and Ta-sro1AOE-R, and the *XbaI*-*Bam*HI-digested PCR product was then ligated into the binary vector pBI121. To generate the Pro*Ubi*:sro1 plasmid for overexpression in wheat, the Ta-sro1 CDS was amplified using primers Ta-sro1WOE-F and Ta-sro1WOE-R and digested with *Bam*HI and *Sac*I for ligation into the

binary vector pTCK303. The binary plasmids were transferred into the *Agrobacterium tumefaciens* strain AGL1 using the freeze-thaw method. Wheat plants were transformed using the shoot apical meristem method (Zhao et al., 2006), while *Arabidopsis* and *At-rbohF* mutant plants were transformed by the floral dip method (Clough and Bent, 1998). The primer sequences used for the above studies are listed in Supplemental Table 3.

### RNA Extraction, cDNA Isolation, RT-PCR, and Real-Time Quantitative RT-PCR

Total RNA was extracted using the TRIzol reagent according to the user manual (Invitrogen). The first strand cDNA was synthesized using an oligo (dT) primer and SuperScript II reverse transcriptase (Invitrogen). The expression of Ta-sro1 transgene in wheat and *Arabidopsis* was confirmed by RT-PCR with the gene-specific primers. Real-time quantitative RT-PCR was performed in an iCycler thermal cycler and detected using the iCycler iQ real-time PCR detection system (Bio-Rad). The reaction mixture comprised 1  $\mu\text{L}$  each of template, forward and reverse primer (0.3  $\mu\text{M}$  each), and FastStart Universal SYBR Green Master (Rox; Roche). The relevant primers are given in Supplemental Table 3. For wheat, *Actin* was used as the internal control and for *Arabidopsis Tubulin*. All reactions were run in triplicate.

### Subcellular Localization of Ta-sro1

To generate Pro35S:sro1-GFP, a *Bam*HI-*Hind*III fragment containing the coding region of Ta-sro1 was amplified by the primers Ta-sro1LF and Ta-sro1LR (Supplemental Table 3) and subcloned into the *Bam*HI and *Hind*III sites of Pro35S:GFP (Lin et al., 2009). The plasmids Pro35S:GFP and Pro35S:sro1-GFP were introduced into *Arabidopsis* mesophyll protoplasts as described (Sheen, 2001). After overnight incubation in the dark, the GFP signal and chlorophyll autofluorescence were examined under a confocal microscope at excitation wavelengths of 488 and 647 nm, respectively (FluoView 1000; Olympus).

### Quantification of Root Length and H<sub>2</sub>O<sub>2</sub> Content

The root length of treated and untreated control seedlings was measured using an image analysis system (WinRHIZO; Régent Instruments). Prior to H<sub>2</sub>O<sub>2</sub> measurement, the wheat roots were excised and immersed for 1 h in PBS solution containing 0.5 mg mL<sup>-1</sup> nitroblue tetrazolium. The staining reaction was stopped by the addition of an excess of 70% (v/v) methanol. The roots were rinsed in fresh 70% (v/v) methanol (Jones et al., 2007) and then observed and photographed under a stereomicroscope (SMZ-800; Nikon). H<sub>2</sub>O<sub>2</sub> content was determined following the protocol of the H<sub>2</sub>O<sub>2</sub> Colorimetric Assay Kit (Beyotime). A 150- to 300-mg sample of fresh tissue was snap-frozen in liquid nitrogen and powdered in a mortar together with 1.5 mL of frozen 5% (w/v) trichloroacetic acid (TCA) and 45 mg of activated charcoal. The homogenate was centrifuged at 18,000g for 10 min at 0°C, and the resulting supernatant filtered through a 45- $\mu\text{m}$  nylon filter. The pH of the filtrate was adjusted to 8.4 with NH<sub>4</sub>OH. After refiltration, a 100- $\mu\text{L}$  aliquot was mixed with 100  $\mu\text{L}$  of the colorimetric reagent, and the reaction mixture was held for 30 min at 30°C. The intensity of the color that developed was spectrophotometrically assessed at 560 nm.

### SCGE

For UV treatment, the *Arabidopsis* and wheat protoplasts were irradiated with UV-C (380  $\mu\text{W cm}^{-2}$ ) for 30 s, 1 min, and 2 min and used directly for SCGE analysis. For H<sub>2</sub>O<sub>2</sub> treatment, 3 or 5 mM H<sub>2</sub>O<sub>2</sub> was added to the suspension and cultured for 12 h before protoplast isolation. These cells were mixed with 0.65% (w/v) low melting point agarose in 1 $\times$  TBE (Tris/Borate/EDTA), and 70  $\mu\text{L}$  was pipetted onto a microscope slide that had previously been coated with 1.5% (w/v) standard agarose and dried in an oven. The preparation was covered with a 22  $\times$  22-mm cover slip and

placed on ice for 5 min. After removing the cover slip, the slide was immersed in 2 M NaCl, 100 mM EDTA, 10 mM Tris, pH 10, 1% (w/v) *N*-lauroyl sarcosine sodium, 6% (v/v) DMSO, and 1% (v/v) Triton X-100 at 4°C for 3 h, washed twice with deionized water, and then transferred to 300 mM NaOH and 1 mM Na<sub>2</sub>EDTA, pH 13, for 40 min. The contents of the slide were subjected to electrophoresis in 1× TBE for 20 min at 2 V/cm and stained in 5 μg mL<sup>-1</sup> ethidium bromide for 10 min. The distribution of fluorescence in the head and tail of the resulting comet of 30 to 50 cells (a measurement that correlates positively with the extent of DNA breakage) (Olive et al., 1990) was assessed. Olive tail moment (OTM) was used to estimate the DNA damage, which represents the product of the percentage of total DNA in the tail (TD) and the tail length (TL), measured from the center of the nucleus toward the end of the tail (OTM = TL × TD). Each assay was performed in triplicate.

### LD<sub>50</sub> for KCN and Respiration Rate in *Arabidopsis* OE Lines

One-week-old seedlings grown on Murashige and Skoog solid medium were transferred to the same medium supplemented with 50 to 600 μM KCN and scored for survival after 14 d. The respiration rate of the roots of *Ta-sro1* OE and VC plants was obtained from 0.1 to 0.2 g (fresh weight) detached root tissue sealed in an airtight cuvette containing phosphate buffer, pH 6.8, and a Clark-type oxygen electrode (Oxy-lab; Hansatech). The capacity of the alternative and cytochrome C pathways was deduced from the difference between respiration in the presence of 0.5 mM KCN or 3 mM SHAM and residual respiration. The latter was measured in the presence of inhibitors of both 3 mM SHAM and 0.5 mM KCN (Bingham and Farrar, 1989).

### Determination of PARP Activity

To express and purify Ta-SRO1, *Ta-sro1*, *Ta-sro1* (V250G), and *Ta-sro1* (T343A) proteins for PARP enzyme activity assay *in vitro*, the *Ta-SRO1* and *Ta-sro1* CDS were amplified using the primer set listed in Supplemental Table 3. *Ta-sro1* (V250G) was generated by site-directed mutagenesis PCR (Ho et al., 1989) using primer pairs *Ta-sro1*PF/V250GR (5'-TGAATCAC-CAGGTTGGCC-3') and V250GF (5'-GGCCAACCTGGTGATTCA-3')/*Ta-sro1*PR. *Ta-sro1* (T343A) was amplified using primer pairs *Ta-sro1*PF/T343AR (5'-TGCACCAATGCCATACACTG-3') and T343AF (5'-CAGTG-TATGGCATTGGTGA-3')/*Ta-sro1*PR. After sequencing, the DNA fragment containing the entire CDS of *Ta-sro1*, *Ta-SRO1*, *Ta-sro1* (V250G), or *Ta-sro1* (T343A) was cloned into the *Bam*HI and *Not*I sites of the prokaryotic expression vector pET24a. The expression cassette for the fusion protein His<sub>6</sub>-tagged *Ta-sro1* was transferred into *Escherichia coli* strain BL21 (DE3), and the subsequently expressed fusion protein was purified from homogenized bacterial cells using a His-tagged protein purification kit (R&D Systems) according to the manufacturer's protocol. The Quick Start Bradford protein assay (Bio-Rad) was used to determine the concentration of purified proteins. Affinity-purified His-fused proteins were used for PARP activity determination using the PARP Universal Colorimetric Assay Kit (Trevigen) following the manufacturer's protocol.

To explore the PARP activity *in vivo*, roots of the wheat cultivars SR3 and JN177 and whole *Arabidopsis* seedlings were snap-frozen in liquid nitrogen and macerated in frozen 1× PARP buffer (Trevigen) containing 0.4 mM phenylmethylsulfonyl fluoride, 0.4 M NaCl, and 1% (w/v) Triton X-100. The homogenate was centrifuged at 18,000g for 10 min at 4°C, and the supernatant was used to determine the total PARP activity using the PARP Universal Colorimetric Assay Kit.

### In Vitro Protein-DNA Binding Assay (Dot Blotting)

Dot blotting was performed as described previously with slight modifications (Tao et al., 2008). *Ta-sro1*-His<sub>6</sub>, *Ta-SRO1*-His<sub>6</sub>, *At*-PARP1-His<sub>6</sub>, and *Ta*-ACO1-His<sub>6</sub> proteins were first blotted onto a polyvinylidene difluoride

membrane. The membrane was blocked with 5% (w/v) nonfat milk in 15 mL of PBST (137 mM NaCl, 2.7 mM KCl, 1.5 mM KH<sub>2</sub>PO<sub>4</sub>, 8 mM Na<sub>2</sub>HPO<sub>4</sub>, pH 7.4, and 0.1% [v/v] Tween 20) at room temperature for 2 h. The blocked membrane was then washed with 15 mL of PBST buffer and preincubated with 15 mL of DNA probe dilution buffer (0.3% BSA and 1% goat serum in PBST) for 1 h. For DNA probe preparation, the calf thymus DNA (Sigma-Aldrich) was covalently labeled using EZ-Link TFPA-PEG<sub>3</sub>-Biotin (Pierce), following the manufacturer's protocol. The biotinylated calf thymus DNA was then added to the probe dilution buffer to a final concentration of 10 μg/mL, followed by incubation with the membrane at room temperature for 1 h. After washing three times with PBST buffer, the membrane was incubated with streptavidin-horseradish peroxidase conjugate (Pierce) in PBST buffer for 30 min. After three washes, the specific protein-DNA binding was detected with ECL Plus Western Blotting Detection Reagents (Pierce), and images were captured by Image Quant 400 (GE Healthcare).

### Determination of Enzyme Activity

To determine CAT, APX, GPX, MDAR, and DHAR activity, ~0.15 g of plant material was homogenized in 1 mL 50 mM KH<sub>2</sub>PO<sub>4</sub>, 0.1 mM EDTA, and 0.3% (w/v) Triton X-100. For GR and NOX, the Triton X-100 was omitted (Grace and Logan, 1996). Total protein was determined using a BCA protein assay kit (Beyotime) following the kit protocols. CAT activity was determined using the procedure of Aebi (1984), monitoring the decrease in absorbance at 240 nm of H<sub>2</sub>O<sub>2</sub> for 1 min at 25°C. APX activity was measured spectrophotometrically at 290 nm according to the method of Nakano and Asada (1981), applying an AsA extinction coefficient of 2.8 mM<sup>-1</sup> cm<sup>-1</sup>. GPX activity was measured according to Benabdellah et al. (2009), in which the decrease in absorbance at 340 nm of a reaction containing 100 mM Tris-HCl, pH 7.4, 0.5 mM EDTA, 1.0 mM Na<sub>2</sub>N<sub>3</sub>, 2 mM H<sub>2</sub>O<sub>2</sub>, 0.25 mM NADPH, 2.25 mM GSH, and 1.0 unit GR was monitored. A molar extinction coefficient for NADPH of 6.22 × 10<sup>3</sup> M<sup>-1</sup> cm<sup>-1</sup> was assumed (Sebastiani et al., 2007). MDAR activity was assayed according to Miyake and Asada (1992) in which the decrease in absorbance at 340 nm of a reaction containing 50 mM HEPES-KOH, pH 7.6, 0.1 mM NADH, 2.5 mM AsA, and 0.3 unit AsA oxidase was monitored. The same molar extinction coefficient for NADPH as above was assumed. DHAR activity was determined according to Dalton et al. (1986) in which the decrease in absorbance at 265 nm of a reaction containing 100 mM HEPES-KOH, pH 7.0, 0.1 mM EDTA, 2.5 mM GSH, and 0.2 mM AsA was monitored. A molar extinction coefficient for AsA of 14 × 10<sup>3</sup> M<sup>-1</sup> cm<sup>-1</sup> was assumed. GR activity was measured based on the method of Grace and Logan (1996) in which the decrease in absorbance at 340 nm of a reaction containing 100 mM Tris-HCl, pH 8.0, 1 mM EDTA, 1 mM GSSG, and 0.2 mM NADPH was monitored. The same molar extinction coefficient for NADPH as above was assumed. NOX activity was measured as for GR activity, omitting GSSG in the reaction (Grace and Logan, 1996).

### Determination of AsA/DHA, GSH/GSSG, and NAD(P)<sup>+</sup>/NAD(P)H Ratios

A modification of the method of Okamura (1980) was used to determine AsA and DHA content. Snap-frozen tissue (0.1 g) was ground in 1 mL of 5% (w/v) TCA and centrifuged at 15,000g for 15 min. Each extract was divided into two, with one part used for the AsA assay, and the other to assess the total AsA content. The DHA concentration was calculated from the difference between the latter and the former. AsA was quantified by first adding 100 μL of 5% (w/v) TCA and 100 μL ethanol to 100 μL of the TCA extract. Then the following were added sequentially: 50 μL of 0.4% (v/v) H<sub>3</sub>PO<sub>4</sub>-ethanol, 100 μL of 0.5% bathophenanthroline-ethanol, and 50 μL of 0.03% (w/v) FeCl<sub>3</sub>-ethanol. The solution was then left to stand for 90 min at 30°C, and the absorbance was read at 534 nm. To assess the total AsA content, a 25-μL aliquot of 1.8 M triethanolamine was added to 100 μL of the TCA extract to neutralize the pH. After incubation at room temperature for 10 min,

50  $\mu$ L of 0.5% (v/v) *N*-ethylmaleimide was added to block any excess reducing reagent, followed by 50  $\mu$ L of 20% (w/v) TCA to obtain a pH of 1 to 2. Thereafter, the protocol followed the one used for AsA determination.

Total GSH was measured by the 5,5'-dithiobis(2-nitrobenzoic acid)-GR recycling assay (Rahman et al., 2006). A 0.1-g plant sample was homogenized in 1 mL of ice-cold 5% (w/v) sulfosalicylic acid and centrifuged at 15,000g for 15 min. The supernatant was neutralized by the addition of 7.5 M triethanolamine. A 200- $\mu$ L aliquot was then used to determine total GSH (GSH+GSSG) content, while 4  $\mu$ L of 2-vinylpyridine was added to a second 200- $\mu$ L aliquot, which was then held at 20°C for 60 min. A volume of 700  $\mu$ L of 0.3 mM NADPH, 100  $\mu$ L of 10 mM 5,5'-dithiobis(2-nitrobenzoic acid), and 150  $\mu$ L of 125 mM NaH<sub>2</sub>PO<sub>4</sub> in 6.3 mM EDTA, pH 6.5, was mixed with 50  $\mu$ L of each of the two aliquots above, and 10  $\mu$ L of 50 units/mL of GR was then added. Both reactions were monitored spectrophotometrically at 412 nm.

NAD(P)<sup>+</sup> and NAD(P)H were quantified by methods adapted from Gibon and Larher (1997). A 0.1-g sample was homogenized in 1 mL of ice-cold 0.1 M HCl (for NAD<sup>+</sup> and NADP<sup>+</sup>) or in 1 mL of 0.1 M NaOH (NADH and NADPH). The homogenates were boiled for 5 min and then cooled on ice and centrifuged (10,000g, 4°C, 10 min). The resulting supernatants were neutralized with either 0.1 M NaOH or 0.1 M HCl (as appropriate) and then recentrifuged (10,000g, 4°C, 10 min). These supernatants were kept on ice during a reduction reaction of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide catalyzed by phenazine methosulfate in the presence of ethanol and either alcohol dehydrogenase (for NAD<sup>+</sup> and NADH) or Glc-6-phosphate and Glc-6-phosphate dehydrogenase (for NADP<sup>+</sup> and NADPH).

#### Determination of ATP and MDA Content

ATP was extracted at 2°C from 250 mg of frozen plant tissues by homogenization with 6 mL of 6% perchloric acid for 30 s in a Potter glass homogenizer. After centrifugation for 15 min at 20,000g at 2°C, the supernatant was neutralized with 5 M ice-cold K<sub>2</sub>CO<sub>3</sub> (~0.36 mL) to pH 7.75 and then precipitated for 1 h in an ice bath. KClO<sub>4</sub> was removed by centrifugation for 15 min at 20,000g at 2°C, and the supernatant was used to determine ATP content immediately with the ENLITEN ATP assay system (Promega) according to the manufacturer's protocol. MDA content was determined according to Liu et al. (2012).

#### Three-Dimensional Model Construction and Analysis

Structural templates for the catalytic domains of the Ta-SRO1, Ta-sro1, At-RCD1, At-PARP1, and At-PARP2 proteins were determined using pGenThreader (Lobley et al., 2009). All recognized templates were assessed as "high" or "certain" by pGenThreader. The initial three-dimensional coordinates of the models were generated with MODELER 9v3 (Fiser et al., 2000). Thereafter, the models were iteratively evaluated with ProQ (Wallner and Elofsson, 2003) and PROCHECK (Laskowski et al., 1993) and locally optimized with ModLoop (Fiser and Sali, 2003), until satisfactory evaluation scores were obtained. The molecular surface electrostatic potential of the resulting models was calculated from the Adaptive Poisson-Boltzmann Solver software plug-in of the Visual Molecular Dynamics program (Humphrey et al., 1996).

#### Accession Numbers

Sequence data from this article can be found in the Arabidopsis Genome Initiative, GenBank/EMBL databases, or the Rice Genome Annotation Project database (<http://rice.plantbiology.msu.edu>) under the following accession numbers: Ta-sro1 (JN202574), Ta-ACO1 (KF014124), Ta-Actin (AB181991), At-RCD1 (At1g32230), At-SRO1 (At2g35510), At-SRO2 (At1g23550), At-SRO3 (At1g70440), At-SRO4 (At3g47720), At-SRO5 (At5g62520), At-AOX1a (At3g22370), At-NDB2 (At4g05020), At-RbohF (At1g64060), At-Tublin (At1g04820), At-ATM (At3g48190), Os-SRO1a (Os10g42710),

Os-SRO1b (Os3g63770), Os-SRO1c (Os3g12820), Os-SRO1d (Os6g13860), and Os-SRO1e (Os4g57640).

#### Supplemental Data

The following materials are available in the online version of this article.

**Supplemental Figure 1.** Variation of Total Root Length and root H<sub>2</sub>O<sub>2</sub> Content in SR3 under Stress Conditions.

**Supplemental Figure 2.** Sequence Alignment of Coding DNA Sequences and Protein Sequences of SRO1 and sro1.

**Supplemental Figure 3.** PCR and qRT-PCR Test of Transgenic Wheat Plants.

**Supplemental Figure 4.** Variation of Total Root Length and Root H<sub>2</sub>O<sub>2</sub> Content in Transgenic Wheat Lines under Stress Conditions.

**Supplemental Figure 5.** In Vitro DNA Binding Activity of Ta-sro1-His<sub>6</sub> and Ta-SRO1-His<sub>6</sub> Proteins.

**Supplemental Figure 6.** Transgenic Test of the Constitutive Expression of Ta-sro1 in Wild-Type *Arabidopsis*.

**Supplemental Figure 7.** Overexpression of Ta-sro1 in the At-rbohF Mutant Background Promoted the Oxidative Stress Tolerance of the Mutant.

**Supplemental Figure 8.** Single-Cell Gel Electrophoresis Analysis of the Protoplasts of JN177 and SR3 Exposed to UV Treatment.

**Supplemental Figure 9.** The ATP Content between Vector Control and Ta-sro1 Overexpression *Arabidopsis* Lines under Normal and Stress Conditions.

**Supplemental Figure 10.** Expression of At-ATM in Vector Control and Ta-sro1 Overexpression *Arabidopsis* Lines.

**Supplemental Figure 11.** The Malondialdehyde Content of Vector Control and Ta-sro1 Overexpression *Arabidopsis* Lines under Salinity Stress.

**Supplemental Table 1.** Oxidative Stress-Related Genes Showing at Least a Twofold Induction or Repression upon Exposure to Abiotic Stress.

**Supplemental Table 2.** Model Evaluations and Templates (PDB Codes) Used for Modeling.

**Supplemental Table 3.** Primers Used for PCR, RT-PCR, and Real-Time RT-PCR Analysis.

**Supplemental Data Set 1.** Alignments Used to Generate the Phylogeny Presented in Figure 2D.

#### ACKNOWLEDGMENTS

This work was supported by funds of the Major Program of the Natural Science Foundation of China (No. 31030053), the National Basic Research 973 Program of China (2012CB114204), and the National Transgenic Project (Grants 2013ZX08002002 and 2013ZX08002003).

#### AUTHOR CONTRIBUTIONS

G.X. designed the experiment. S.T.L. performed most of the experiments. S.W.L. and Mei W. performed some key experiments. T.W. performed three-dimensional model construction and analysis. C.M. characterized transgenic plants. Meng W. performed the SCGE experiment. G.X., S.T.L., S.W.L., and Mei W. wrote the article.

Received September 18, 2013; revised November 26, 2013; accepted December 26, 2013; published January 17, 2014.

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