The Maize Sulfite Reductase Is Involved in Cold and Oxidative Stress Responses

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Sulfite reductase (SiR) functions in sulfate assimilation pathway. However, whether it is involved in stress response in crops is largely unknown. Here, the SiR ortholog from Zea mays (ZmSiR) was characterized. The recombinant ZmSiR protein was purified from E. coli. It exhibited sulfite-dependent activity and had strong affinity for sulfite. ZmSiR transcripts were markedly up-regulated by cold and methyl viologen (MV) treatments. Overexpression of ZmSiR complemented growth retardation phenotype of Arabidopsis atsir mutant. ZmSiR-overexpressing Arabidopsis plants were tolerant to severe SO2 stress and rescued the susceptible phenotype of the atsir. ZmSiR knock-down transgenic maize plants with 60% residual transcripts were more susceptible to cold or oxidative stress than wild-type. The severe damage phenotypes of the ZmSiR-compromised maize plants were accompanied by increases of sulfite and H2O2 accumulations, but less amounts of GSH. The qPCR analysis revealed that there was significantly altered expression of several key sulfur metabolism-related genes in ZmSiR-impaired maize lines under cold or MV stress. Particularly, ZmAPR2 expression was significantly elevated, suggesting that toxic sulfite accumulation in ZmSiR-impaired plants could be attributable to the reduced SiR coupled to increased ZmAPR2 expression. Together, our results indicate that ZmSiR is involved in cold and oxidative stress tolerance possibly by modulating sulfite reduction, GSH-dependent H2O2 scavenging, and sulfur-metabolism related gene expression. ZmSiR could be exploited for engineering environmental stress-tolerant varieties in molecular breeding of maize.

Keywords: maize, sulfite reductase, cold stress, oxidative stress, glutathione

INTRODUCTION

In higher plants, the assimilatory sulfate reduction is an essential metabolic process in which a great number of sulfur-containing amino acids, sulfolipids and coenzymes are synthesized (Leustek and Saito, 1999; Leustek et al., 2000; Lewandowska and Sirko, 2008). This process involves several successive enzymatic reactions that occur in chloroplasts. Firstly, sulfate is transported to chloroplasts, where it is adenylated by ATP sulfate adenyllytransferase (APS) (Cao et al., 2013). Then, the product adenosine-phosphosulfate is reduced to sulfite by APS reductases (APRs). And
then, reduction of the toxic sulfite to sulfide is catalyzed by sulfite reductase (SiR). Finally, the sulfide is further incorporated into cysteine by O-acetyl-serlyases that can be used for synthesizing other sulfur-containing compounds such as glutathione (Nakayama et al., 2000).

Plant SiR is a kind of soluble protein that contains one iron-sulfur cluster (4Fe-4S) and one siroheme group. It is localized to plastids and catalyzes reduction of sulfite to sulfide using reduced ferredoxin (Fd) as a physiological electron donor (Nakayama et al., 2000). It was evidenced that SiR protected Arabidopsis or tomato plants against sulfite/SO$_2$ toxicity via sulfite reduction-dependent pathway (Yarmolinsky et al., 2013). Thus, SiR has been thought to be a bottleneck for assimilatory sulfate reduction in plants (Khan et al., 2010). In addition to reductive assimilation, plant SiR plays an important role in normal growth and development. For example, Nicotiana benthamiana SiR (NbSiR) protein has been evidenced to act as a chloroplast (cp) nucleoid protein with cp DNA-compacting and transcription-regulatory activities (Sekine et al., 2007; Kang et al., 2010). Furthermore, the conserved C-terminally encoded peptides of the SiR are important to localize to the cp nucleoids in plants (Kobayashi et al., 2016). NbSiR silencing differentially affected expression of plastid-encoded genes and ultimately resulted in chloroplast ablation (Kang et al., 2010), indicating that plant SiR functions in cp-nucleoid metabolism, plastid gene expression, and chloroplast development. In addition, it was reported that Arabidopsis plants with impaired SiR expression exhibited dramatically reduced biomass and marked metabolism disturbances (Khan et al., 2010). Also, tomato plants with lowered SiR activity showed clearly reduced biomass and promoted early leaf senescence (Yarmolinsky et al., 2014). These results indicate that SiR is essential for plant growth and development.

Although new insights in biological functions of SiR were reported in model plants, the knowledge of SiR function and regulation from crops is very limited. Environmental stresses such as drought, cold, and high salinity cause oxidative stress by provoking cellular redox imbalances, thus seriously affecting growth, development, and biomass of crop plants (Gechev et al., 2006). To date, information about biochemical properties and biological function of maize SiR is scarce. In this study, we characterize enzymatic properties of the maize SiR and its function in cold and oxidative stresses in maize.

**MATERIALS AND METHODS**

**Plant Material and Growth Conditions**

The maize inbred line Zheng58-1 and Arabidopsis thaliana ecotype Columbia (Col-0) were used for this study. For Arabidopsis, the seeds were surface sterilized and then germinated on plates containing one-half-strength Murashige and Skoog (MS) medium. After 7 days, the seedlings were transferred to sterilized low-nutrient soil to obtain fully grown plants. For maize, the seeds were germinated on moist filter papers in the sterilized low-nutrient soil to obtain fully grown plants. For on plates containing one-half-strength Murashige and Skoog Arabidopsis ecotype Columbia (Col-0) were used for this study. For Arabidopsis thaliana The maize inbred line Zheng58-1 and were grown in a growth room at approximately 23°C, 60–70% relative humidity, a photoperiod of 18 h/6 h (day/night) and light intensity of 200 µmol m$^{-2}$ s$^{-1}$, as described previously (Xia et al., 2013).

**Stress Treatments on Maize Plants**

For oxidative stress, 2-week-old Zheng 58-1 plants in soil were sprayed with 20 µM of MV. For cold stress, the same age maize plants in soil were transferred to grown at 4°C in the growth chamber. During 48 h of both types 58st NL C-Terminal Region types of stress treatments, leaf samples were harvested at indicated time points (0, 1, 3, 6, 12, 24, and 48 h) for gene expression analyses.

**Overexpression, Purification, Immunoblot Detection, and Kinetic Analysis of Recombinant ZmSiR**

The ZmSiR full-length cDNA fragment was amplified and cloned into the vector pET-30a (Novagen, United States) with NdeI/XhoI sites (Supplementary Table S1). The resulting expression construct pET-ZmSiR was transformed into E. coli BL21 (DE3) pLysS cells, which could produce a recombinant protein with a 6× histidine tag at its N-terminus by 0.5 mM of isopropylthio-β-galactoside (IPTG) induction. The cells were harvested by centrifugation at 8,000 × g for 10 min and resuspended in a solution containing 0.1 M Tris acetate (pH 7.6), 0.5 M sucrose, 0.1 mM DTT, 0.5 mM EDTA, and Protease inhibitor cocktail, and then sonication was conducted using an ultra-sonicator (450 Digital Sonifier, United States) on ice with 50% pulse amplitude. After that, the bacterial fractions containing ZmSiR-His were centrifuged (8,000 × g, 20 min) and the precipitations were solubilized at 4°C overnight in a dissolution buffer [0.1 M potassium phosphate buffer (pH 7.4), 10 mM β-mercaptoethanol, 0.5 M NaCl, 20% (w/v) glycerol, 1, 1.0% (w/v) CHAPS, and 0.5% (v/v) NP-40]. The purification of ZmSiR-His protein was carried out using a Ni$^{2+}$-affinity purification kit (Sangon, Co., Shanghai, China) as previously described (Xia et al., 2012). Briefly, the solubilized fraction was loaded onto a pre-equilibrated Ni$^{2+}$-affinity column and bound proteins were washed with a washing buffer containing 0.1 M potassium phosphate buffer (pH 7.4), 10 mM β-mercaptoethanol, 0.5 M NaCl, 20% (w/v) glycerol, and 30 mM imidazole. The purified protein was obtained using an elution buffer that had same composition as the washing buffer, except that 30 mM imidazole was replaced by 20 mM imidazole. The eluted fraction containing the ZmSiR-His protein was collected and dialyzed at 4°C against 100 mM potassium phosphate buffer (pH 7.4) containing 20% (v/v) glycerol.

For immunoblot detection of the recombinant SiR, the cell extracts or purified proteins were separated using 12% SDS-Tris-glycine polyacrylamide gel electrophoresis (SDS-PAGE) and transferred onto polyvinylidene difluoride (PVDF) membranes. The membranes were blocked and thereafter blotted with a commercial His-tagged monoclonal antibody (CWBiO, Co., Beijing, China) for 3 h at a 1:3000 dilution. After extensive washing, the bound primary antibody was detected with a horseradish peroxidase (HRP)-conjugated
goat anti-mouse IgG secondary antibody (CWBIO Co., Beijing, China). The HRP activity was visualized using a 3,3′-diaminobenzidine (DAB) development kit (CWBIO, Co., Beijing, China) as described by us (Xia et al., 2012).

Kinetic analysis of ZmSiR was conducted using a saturating concentration of NADPH (400 µM) and varying sulfite concentrations (from 2.5 to 400 µM). The kinetic parameters (Km and Vmax) were calculated as described previously (Xia et al., 2012). For the assays, three replicates were conducted for each test sample and the experiment was repeated three times.

**Verification of SiR Arabidopsis Mutants**

The AtSiR T-DNA insertion mutant (SALK_047219) (named atsir-s047) seeds were obtained from the ABRC collection center (The Ohio State University, Columbus, OH, United States). Homozygous mutants were identified by PCR from genomic DNA using forward primer P1, T-DNA left border primer LBb1, and AtSiR (Accession No. At5g04590) gene-specific reverse primer P2 (Supplementary Table S1), and analyzed further by DNA sequencing to confirm the insertions of the T-DNA in the gene. The transcript levels of AtSiR in the Col-0 and mutants were determined by quantitative PCR.

**Construction of atsir/35S-ZmSiR Transgenic Arabidopsis Plants and SO2 Stress Phenotype Analysis**

The full-length ZmSiR cDNA was amplified by PCR using corresponding primer pairs (Supplementary Table S1), and was subsequently into the binary vector pART27 (Xia et al., 2012).
ZmSiR transgenic plants were exposed to 6 ppm SO\(_2\) analysis.

Homozygous T3 Arabidopsis lines were used for phenotypic analysis. Five-weeks old wild-type, atsir mutant, and atsir/35S-ZmSiR transgenic plants were exposed to 6 ppm SO\(_2\), or to filtered pollutant-free air (control) for 2 h in fumigation chambers. The gas was released through a tube from a cylinder and kept constant by a SO\(_2\) monitor in the chamber as described by us (Xia et al., 2012). After treatments, the plants were transferred to the growth room for recovery. Three replicates each consisting of three pots of seedlings from each line were included for both SO\(_2\) treatment and non-treated controls. The whole experiment was repeated three times.

**Expression, purification, and kinetic analysis of recombinant maize sulfite reductase.** (A) Purification of histidine-tagged maize SiR. The histidine-tagged ZmSiR (with a predicted molecular mass of 70 kDa) was expressed in bacterial cells induced by 0.5 mM isopropyl-β-D-thiogalactopyranosid (IPTG) (lane 2, indicated by white arrowhead). The overexpressed protein was purified using Ni\(^{2+}\)-affinity chromatography (lane 3). The overexpressed and purified SiR proteins were blotted with an anti-histidine monoclonal antibody and the blotted target bands were visualized using a 3,3′-diaminobenzidine (DAB) development kit based on the HRP activity (lane 4 and lane 5; signals marked by an arrowhead). The overexpressed protein was purified using Ni\(^{2+}\)-affinity chromatography (lane 3). The overexpressed and purified SiR proteins were blotted with an anti-histidine monoclonal antibody and the blotted target bands were visualized using a 3,3′-diaminobenzidine (DAB) development kit based on the HRP activity (lane 4 and lane 5; signals marked by an arrowhead). The molecular mass markers (kDa, lane 1) are shown on the left. (B) Steady-state kinetics of recombinant ZmSiR with sulfite. Double-reciprocal presentation (Lineweaver–Burk plot) of enzyme rate was conducted using varying concentrations of sulfite (2.5, 5, 10, 25, 50, 100, 200, and 400 µM) and constant 400 µM NADPH. The reaction was initiated with 1.0 µg of purified recombinant ZmSiR.

**Generation and Molecular Verification of ZmSiR-Compromised Transgenic Maize Lines**

For the ZmSO-compromised construct, a 531-bp-length PCR product of ZmSiR cDNA was amplified using primers ZmSiR-F2 and ZmSiR-R2 (Supplementary Table S1) and introduced as sense and anti-sense into the binary vector pFGC491 with BamHI and XbaI, or NcoI and SmaI restriction sites, resulting in the transformation construct pFGC491-35S: ZmSiR-RNAi. The recombinant plasmid was introduced into the A. tumefaciens strain LBA4404, which was used to transform maize. Transformation of maize inbred line Zheng58-1 immature embryos was conducted as described by us previously (Xia et al., 2015). Transgenic maize seedlings with Basta resistance were screened and confirmed by PCR as described previously (Xia et al., 2015). The number of T-DNA inserts in PCR-positive plantlets was assessed by segregation of T\(_0\) offspring on selective medium. Several T\(_0\) seedlings of each one-copy line were grown in the field and their seeds were germinated and used for stress assays. The transcript levels of ZmSiR-compromised transgenic lines were evaluated by RT-qPCR as described below.

**Analysis of ZmSiR-Compromised Maize Lines for Oxidative or Cold Stress Tolerance**

For oxidative stress, 3-week-old maize WT and ZmSiR-compromised lines (S2, S5, and S6) were sprayed with 20 µM of MV. Three replicates each consisting of two pots of seedlings from each line were included for both MV treatment and H\(_2\)O\(_2\)-treated controls. For cold stress, 12-day-old WT and both ZmSiR-compromised lines (S5 and S6) were transferred to grow hydroponically in MS liquid medium at 4°C (Cold) or 23°C (Control) for 3 days. After treatments, phenotypes of various types of plants were photographed. Total chlorophyll content in both WT and RNAi plants was determined as described by Arnon (1949). In brief, leaves were sampled and ground in 80% acetone and the homogenate was centrifuged. Then, absorbance of the supernatant was measured at 645 and 663 nm using a spectrophotometer (Hitachi U2000, Japan). Finally, the relative remaining chlorophyll (%) of the WT and both RNAi maize lines was determined. The experiments were repeated three times each.

**Histochemical and Quantified Detection of H\(_2\)O\(_2\) Accumulation**

The histochemical detection of H\(_2\)O\(_2\) from MV-treated maize seedlings with 3, 3′-diaminobenzidine (DAB) was performed according to our previous method (Xia et al., 2013). After staining, the leaf disks were rinsed in 80% (v/v) ethanol for 10 min at 70°C, mounted in lactic acid:phenol:water (1:1:1, v/v/v), and photographed directly using a digital camera. H\(_2\)O\(_2\) content in various genotypes of maize plants was assayed according to our published method (Xia et al., 2012).
**Determination of SiR Activity, Sulfite, and Glutathione Contents**

Sulfite reductase activity assay was determined as described by Khan et al. (2010). Sulfite and reduced glutathione (GSH) contents of leaves were measured using ion-exchange chromatograph system as described by us (Wang et al., 2016). For each assay, three replicates were done for each test sample and these experiments were repeated three times.

**Quantitative Real-Time PCR (RT-qPCR)**

Total RNA extraction was conducted as described previously (Xia et al., 2012). The first-strand cDNA was synthesized from 1 µg of total RNA using the SuperScript™ RT kit (Thermo Scientific, United States) in a 10 µL-reaction volume according to the manufacturer’s recommendations.qPCR was performed in 96-well white plates using an IQ5 Real Time PCR (Bio-Rad, Hercules, CA, United States) with three biological replicates and three technical replicates. The 20 µL reaction mixture consisted of 1 µL cDNA diluted 10-fold, 10 µL master mix (SYBR Green Supermix, Thermo Scientific, United States), and 0.5 µM of each gene-specific primer (Supplementary Table S1). The qPCR procedures were performed according to our detailed descriptions (Su et al., 2017). Data were normalized to the corresponding reference genes transcripts (AtActin2 and AtTubulin for Arabidopsis, and ZmActin2 and ZmTubulin for maize) and the transcript levels of genes were calculated according to the $2^{-\Delta\Delta Ct}$ method (Livak and Schmittgen, 2001). All qPCR experiments followed the MIQE guidelines (Bustin et al., 2009).

**Statistical Analysis**

The data in this study are presented as the mean ± SE of three replicates and were analyzed by a simple variance analysis (ANOVA). The differences between the means were compared by Student’s $t$-test at $P < 0.05$. SPSS statistical software (version 17.0) was used to perform the analyses.

**Accession Numbers**

Sequence data in this study can be found in the GenBank under the following Accession Nos. ZmSO, FJ436404; ZmSiR, NP_001105302; ZmAPR2, GRMZM2G087254; ZmCys2, LOC109939155; ZmSQD1, LOC100282424; ZmSULTR1;1, GRMZM2G159632; ZmSULTR3;1, GRMZM2G154211; ZmGSH1, LOC103648180; ZmGSH2, LOC542055.

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**FIGURE 3** | Overexpression of ZmSiR in the atsr mutant Arabidopsis and its response to toxic levels of SO$_2$ exposure. (A) Toxic effect of SO$_2$ on atsr-s047 and transgenic Arabidopsis plants. Five-week-old WT, atsr-s047 and both atsr/35S-ZmSiR lines (#3 and #7) were exposed to 6 ppm of SO$_2$ for 2 h and examined 4 days later. (B) Relative damage levels in the atsr-s047 and both transgenic Arabidopsis lines after SO$_2$ exposure. (C) Relative residual chlorophyll content in the wild-type, atsr-s047 and atsr/35S-ZmSiR Arabidopsis plants after SO$_2$ exposure. Data are means of three independent repetitive experiments ($\pm$ SE), $n$ = 20. **$t$-test, with $P < 0.01$ and *$t$-test, with $P < 0.05$.**
RESULTS

Molecular Characterization of Maize Sulfite Reductase Gene (ZmSiR)

At the start of this work, total RNAs from 2-week-old maize plants, which had been treated with 20 µM methyl viologen (MV) for 12 h, were used as samples for transcriptome analysis to identify genes responsive to oxidative stress. A gene encoding sulfite reductase in maize (named ZmSiR) with nearly 10-fold induction was observed (data not shown). The full-length cDNA of ZmSiR from maize was obtained by RT-PCR. The cDNA sequence of ZmSiR is 2370 bp long, which consists of a 94 bp 5′-untranslated region, a 368 bp 3′-untranslated region, and a 1908 bp ORF. Sequence comparison between gDNA and cDNA revealed that the entire ZmSiR genomic sequence contains seven introns and eight exons that encode a 635 amino acid polypeptide (Figure 1A). Amino acid sequence comparisons have revealed that ZmSiR exhibits high identity to counterpart proteins from Arabidopsis thaliana (71% identity), Solanum lycopersicum (65% identity), Ricinus communis (67% identity), Nicotiana benthamiana (66% identity), Glycine max (68% identity), Oryza sativa (91% identity), Brachypodium distachyon (89% identity), Hordeum vulgare (89% identity), and Sorghum bicolor (97% identity) (Supplementary Table S2).

A phylogenetic tree was established based on SiR protein sequences available in GenBank from 10 plant species including Arabidopsis, tobacco, tomato, soybean, castor, rice, barley, wild wheat, sorghum, and maize (Figure 1B). As shown in Figure 1B, these SiRs were clustered into two distinct groups, monocot and dicot SiR groups. In the monocot SiRs, the SiR from maize showed higher identities with sorghum, and thus was clustered into the same isoform subgroup. On the other hand, the SiRs from rice, barley and wild wheat formed another subgroup. In the dicot SiRs, the SiRs from tobacco and tomato were clustered into the same isoform subgroup, while the SiRs from other three species Arabidopsis, soybean and castor formed another subgroup (Figure 1B). These results clearly demonstrated that ZmSiR is highly similar to the known SiR proteins from Arabidopsis and other plant species, thus could be identified as an ortholog of SiR.

Transcript Profiles of ZmSiR in Response to Oxidative or Cold Stress

Time-course analysis of ZmSiR transcript levels in maize plants in response to oxidative stress was performed by qPCR. As shown...
in Figure 1C, the transcript level of ZmSiR was increased rapidly at 3 h, and reached a maximal level at 12 h (about eightfold increase in transcripts) during 48 h of MV treatment (Figure 1C). This suggests that ZmSiR could be involved in oxidative stress response.

Also, transcriptional response of ZmSiR to cold stress in maize plants was examined by qPCR. The expression level of ZmSiR displayed a significant increase at 6 h of cold stress, and reached a maximal level at 12 h (about sixfold increase in transcripts), and then gradually decreased, finally maintained to a relatively high level during 48 h of the treatment (Figure 1D). This result indicates that ZmSiR is also responsive to cold stress.

**Sulfite Reductase Activity of ZmSiR in vitro**

To ascertain whether the cloned ZmSiR encodes a functional sulfite reductase, we have inducibly over-expressed the hexahistidine (His)-tagged ZmSiR protein in *E. coli* and purified it using Ni²⁺-affinity chromatography (Figure 2A; left panel).

The purified ZmSiR protein had a MW of 70 kDa as shown by Western blot assays. This mass was in agreement with our prediction (Figure 2A; right panel). Subsequently, biochemical assays were conducted to analyze its kinetics, in which the recombinant ZmSiR protein showed a sulfite-dependent activity when NADPH acted as an electron acceptor. In the assay, the Michaelis constant $K_m$ for sulfite was calculated to be 5.32 ± 0.26 µM, and the maximum velocity $V_{max}$ for sulfite was calculated to be 10.84 ± 0.12 µM min⁻¹ (Figure 2B). Similar kinetic parameters were also produced in our independent biological replicates. These results suggested that ZmSiR has sulfite reductase activity in vitro.

**ZmSiR Could Complement Growth Retardation Phenotypes of Arabidopsis SiR Mutant**

To clarify whether ZmSiR gene is a functional ortholog of Arabidopsis SiR, one T-DNA insertion line SALK_047219 (named...
atsir-s047) was identified in the ABRC seed stock center. In this line, the T-DNA was inserted in the promoter region of AtSiR (Supplementary Figure S1A). Noticeably, the T-DNA lines located in the gene coding region were embryo lethal (Khan et al., 2010). The T-DNA insertion position is illustrated in Supplementary Figure S1A, and homozygous and heterozygous mutants were verified using AtSiR gene-specific and T-DNA border primers. In accordance with previous findings by Khan et al. (2010), the atsir-s047 homozygous line also showed retarded growth phenotypes (Supplementary Figure S1E). qPCR assays revealed that a decrease of 50% approximately in mRNA abundance was detected compared with wild-type plants of the same age (Supplementary Figure S1B). Accordingly, the SiR activity also was lowered to about 50% of the wild-type level (Supplementary Figure S1C). To verify whether ZmSiR could complement growth retardation phenotypes of the atsir-s047 mutant, the construct harboring the complete ORF of ZmSiR was transformed into atsir-s047 mutant, and two homozygous transgenic Arabidopsis lines (atsir/ 35S-ZmSiR#3 and 7) were used for further analysis. The enzymatic activity determination showed that these transgenic lines had higher SiR activities than the WT, except for the mutant atsir-s047 (Supplementary Figure S1D). Under normal conditions, the atsir-s047 plants showed clearly retarded growth phenotypes compared to the WT, whereas both transgenic lines (atsir/ 35S-ZmSiR#3 and 7) grew normally, similar to the WT (Supplementary Figure S1E). Further biomass determination showed that both complemented Arabidopsis lines had increased by 30–40% in dry weight per plant, compared with the WT (Supplementary Figure S1F). These results suggest that ZmSiR could effectively complement growth retardation phenotypes of the atsr mutant Arabidopsis, which was due to deficiency of SiR activity.

ZmSiR Could Rescue Phenotypes of Arabidopsis SiR Mutant Plants Susceptible to Toxic SO$_2$ Stress

A previous study has evidenced that SiR-impaired Arabidopsis plants are susceptible to excess SO$_2$ because of loss of sulfite reduction-dependent sulfite detoxifying capability (Yarmolinsky et al., 2013). In view of this, it was interesting to examine whether ZmSiR could rescue phenotypes of Arabidopsis SiR mutant susceptible to toxic SO$_2$. The WT, atsir-s047, and both

![FIGURE 6](image-url) | Effect of ZmSiR suppression on transcripts of sulfur metabolism-related genes under oxidative stress. Leaf samples from 10-day-old maize seedlings from wild-type and ZmSiR-compromised lines (S5 and S6) were harvested after 24 h of MV (20 µM) spraying, and then transcript levels of genes encoding sulfate transporters (ZmSULTR1;1 and ZmSULTR3;1), ATPS reductase (ZmAPR2), sulfite oxidase (ZmSO), sulfolipid biosynthesis enzyme (ZmSQD1), cysteine synthetase (ZmCys2), and glutathione synthetase (ZmGSH1 and ZmGSH2) were detected by qPCR. The mRNA levels of these genes were normalized to the transcripts of ZmActin2 or ZmTubulin in the same samples. For each assay, the expression level of WT under control conditions was taken as 1.0, and data represented mean ± SE of three biological replicates. **t-test, with $P < 0.01$; *t-test, with $P < 0.05$. 
atsir/35S-ZmSiR lines (#3 and #7) were exposed to 6 ppm of SO$_2$ for 2 h. After 4 days recovery, three types of Arabidopsis plants all displayed severe leaf injuries (Figure 3A). Comparatively, both ZmSiR-transgenic Arabidopsis lines showed relatively less necrosis and chlorophyll bleaching than the atsir-s047 and wild-type plants (Figure 3A). The leaf damage levels of both atsir/35S-ZmSiR lines were much lower than that in the atsir-s047 line (Figure 3B). Also, the index of residual chlorophyll among these types of plants showed similar changes (Figure 3C). These results have clearly demonstrated that ZmSiR-overexpressing transgenic plants, as opposed to atsir mutant plants, were tolerant to severe SO$_2$ stress and could effectively rescue the susceptible phenotypes of the atsir mutant Arabidopsis.

**Generation of ZmSiR-Compromised Transgenic Maize Lines and Their Response to Oxidative Stress**

To further explore ZmSiR functions, we employed the transgenic approach to silence the gene in maize. A total of 10 independent maize transgenic lines were generated, and the transgenes were confirmed by PCR (data not shown), six of which (S1–S6) were characterized in more detail. As shown in Figure 4A, the decreases in the ZmSiR transcript levels were observed among the six lines by qPCR. Compared with the non-transgenic control (WT), the transcript levels among the six ZmSiR RNAi lines decreased by 20–45% (Figure 4A). Among the six lines, three lines (S2, S5, and S6) with sufficient seeds were chosen for further experiments.

Seeds of the three RNAi maize lines were directly sown in soil for the MV treatment experiment. Under normal conditions, these three RNAi lines showed slight growth retardation at the four-leaf seedlings stage compared to WT plants (Figure 4B). Four days after MV treatment, the RNAi transgenic lines showed relative higher necrosis and wilting than the WT plants (Figure 4B). Accordingly, remaining chlorophyll content in these three RNAi lines was significantly lower (35, 45, and 41% decreases, respectively) than that in the WT plants (16% decrease) (Figure 4C). This result showed that impairment of ZmSiR in maize plants also decreased tolerance to MV-induced oxidative stress.

**Accumulations of Reactive Oxygen Species and S-Metabolites in ZmSiR-Compromised Maize Lines Under Oxidative Stress**

We further detected reactive oxygen species (ROS) accumulations in the WT and RNAi lines during oxidative stress. Histochemical detection of H$_2$O$_2$ in MV-treated and control
Changes in Sulfur-Metabolism Related Gene Transcripts in ZmSiR-Compromised Maize Lines Under Oxidative Stress

The transcripts of S-metabolism related genes encoding sulfate transporters (ZmSULTR1; 1 and ZmSULTR3; 1), APS reductase (ZmAPR2), sulfite oxidase (ZmSO), sulfolipid biosynthesis enzyme (ZmSQQD1), cysteine synthetase (ZmCyS2), and glutathione synthetase (ZmGSH1 and ZmGSH2) were examined under oxidative stress in WT and both RNAi maize lines by qPCR. After 24 h of MV exposure, transcripts of all the genes except for ZmAPR2 in both RNAi plants displayed significant decreases when compared to the WT (Figure 6). In contrast, elevated expression of the ZmAPR2 was quite clear in both RNAi maize lines compared with that in the WT. In addition, under control conditions, transcriptional expression of ZmAPR2 was significantly up-regulated, while expression of both ZmGSH1 and ZmGSH2 was significantly decreased (Figure 6).

Response of ZmSiR-Compromised Maize Lines Under Cold Stress

Due to markedly transcriptional response of the ZmSiR to low temperature (Figure 1D), we tried to examine the performance of ZmSiR-compromised maize plants under cold stress. The hydroponic maize seedlings from WT and both ZmSiR-compromised lines were transferred to grow at 4°C (Cold) or 23°C (Control). After 3 days, both RNAi maize seedlings showed remarkably more leaf yellowing and necrosis than the WT (Figure 7A). In accordance with this phenotype, remaining chlorophyll in both RNAi lines was significantly lower than that in the WT (Figure 7B). This evidenced that knock-down of ZmSiR in maize plants also decreased cold stress tolerance.

H2O2 levels were quantitatively determined after 24-h cold stress. As shown in Figure 7C, the H2O2 levels in both ZmSiR-compromised lines increased by 87% averagely, while the WT only increased by 55% (Figure 7C). Furthermore, sulfite and GSH contents were determined in both RNAi and WT maize plants under cold stress. As shown in Figures 7D,E, under cold stress, sulfite levels in both RNAi maize lines increased by 100% averagely, whereas WT plants only increased by 65% (Figure 7D). Accordingly, under cold stress GSH levels in both RNAi maize lines significantly decreased, but not in the WT (Figure 7E). These results suggested that impairment of ZmSiR also resulted in insufficient GSH levels, and thus reduced GSH-dependent ROS scavenging capability in maize plants under cold stress. Additionally, the expression of four key S-metabolism related genes ZmSO, ZmAPR2, ZmGSH1, and ZmGSH2 was examined in ZmSiR-compromised maize lines under cold stress by qPCR. After 24 h of cold stress, transcripts of the three genes ZmSO, ZmGSH1, and ZmGSH2 displayed significant decreases, whereas ZmAPR2 expression increased significantly in both RNAi plants when compared to the WT (Supplementary Figure S2). Moreover, transcriptional expression of ZmAPR2 was significantly elevated even under control conditions (Supplementary Figure S2B).

DISCUSSION

As a key enzyme in the reductive sulfate assimilation, SiR plays important roles in sulfite detoxification, growth and development, and senescence control in plants (Khan et al., 2010; Yarmolinsky et al., 2013, 2014). In the present work, we report molecular and biochemical properties, and physiological functions of SiR from maize using several complementary approaches. Our genetic evidence suggests that ZmSiR participates in cold or oxidative stress tolerance through modulating sulfite homeostasis and GSH-mediated ROS scavenging. To our knowledge, this is the first sulfite reductase gene from monocots to be functionally characterized.

SiR exists in a single copy in Arabidopsis and tomato, and in two copies in poplar and rice (Bork et al., 1998; Kopriva, 2006; Brychkova et al., 2012). In maize, ZmSiR was found to exist only as a single gene on the 6th chromosome. The established phylogenetic tree based on SiR protein sequences from 10 plant species clearly demonstrated that ZmSiR is highly similar to the known plant SiRs, suggesting SiR to be evolutionarily conserved in plant species. In addition to sulfite detoxification, SiR functions in growth and development, and leaf senescence regulation (Khan et al., 2010; Yarmolinsky et al., 2013, 2014). In this study, ZmSiR confers cold and oxidative stress tolerance in maize (Figures 4, 7). Together, these results indicate that plant SiR is functionally divergent.

The recombinant ZmSiR protein exhibited a sulfite-dependent sulfite reductase activity (Figure 2). The Km of ZmSiR for sulfite was calculated to be 5.32 μM, which is much lower than the
to those in the WT plants (Figure 6 and Supplementary Figure S2). Moreover, sulfite content significantly increased in the leaves of ZmSiR knock-down maize lines during cold or oxidative stress (Figures 5C, 7D). These observations suggest that toxic sulfite accumulation in ZmSiR-impaired maize plants could be attributable to the reduced SiR coupled to increased ZmAPR2 expression. In accordance with this conclusion, it was reported that constitutive expression of a bacterial APR in Arabidopsis resulted in more than threefold sulfite accumulation and showed chlorotic symptoms of toxicity, suggesting that APR over-expression induced massive de-regulation of primary sulfur metabolism (Tsakraklides et al., 2002). Together, ZmSiR is indispensable for maintaining sulfite homeostasis in maize during cold or oxidative stress.

In summary, ZmSiR has a sulfite-dependent SiR activity with higher affinity toward sulfite. Down-regulation of ZmSiR caused more sulfite and H₂O₂ accumulations, but less GSH levels, and high sensitivity to cold and oxidative stresses in maize. ZmSiR can protect plants from cold and oxidative stresses, and has a potential for engineering environmental stress-tolerant maize varieties in molecular breeding. Future work will be interesting to explore the mechanisms by which ZmSiR is involved in cold or oxidative stress signaling using ZmSiR mutant lines in maize.

AUTHOR CONTRIBUTIONS
ZX designed the research and wrote the manuscript. ZX, MW, and ZXu performed the research and conducted data analyses.

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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2018.01680/full#supplementary-material

FIGURE S1 | Genetic complementation of growth defect of the atsar mutant Arabidopsis. (A) Structure of the AtSiR locus with the T-DNA insertion site in the Salk_047219 (atsar_s047) mutant. The insertion site is marked by a white arrow, exons are indicated as white boxes, and untranslated regions by black boxes. P1, forward primer; P2, reverse primer; LBB1, primer specific to the T-DNA left border. (B) Transcript levels of AtSiR in the wild-type and atsar_s047 determined by qPCR. (C) Relative SiR activity in leaf extracts from wild-type and atsar_s047 plants measured by kinetic assays. (D) Relative activity levels of ZmSiR in the atsar-s047 and transgenic Arabidopsis lines. SiR activity levels of three homozygous atsar/3SS-ZmSiR transgenic lines (named atsar/3SS-ZmSiR #1, #3, and #7), along with the atsar-s047 mutant and wild-type (WT), were determined by spectrophotometry. (E) Representative growth phenotypes of 2-week-old seedlings from atsar-s047 and transgenic Arabidopsis lines. (F) Dry weight of 2-week-old individual seedlings from atsar-s047 and transgenic Arabidopsis lines. In the Figures (B–F), asterisks on the histograms indicate significance of the
difference from the corresponding control values determined by Student’s t-test (**P < 0.01), t-test, with P < 0.05.

FIGURE S2 | Effect of ZmSiR suppression on transcript levels of several key sulfur metabolism-related genes under cold stress. Leaf samples from 12-day-old maize seedlings from wild-type and ZmSiR-compromised lines (S6 and S6) were harvested after 24 h of 4°C (Cold) or 23°C (Control) treatment, and transcript levels of ZmSO (A), ZmAPR2 (B), ZmGSH1 (C), and ZmGSH2 (D) were detected by qPCR as described above. Data represented mean ± SE of three biological replicates. **t-test, with P < 0.01; t-test, with P < 0.05.

TABLE S1 | PCR primers used in this study.

TABLE S2 | Percentages of amino acid sequence identities of SfPs between maize and other plant species.

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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