The **LIKE SEX FOUR2** regulates root development by modulating reactive oxygen species homeostasis in *Arabidopsis*

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Maintaining reactive oxygen species (ROS) homeostasis plays a central role in plants, and is also critical for plant root development. Threshold levels of ROS act as signals for elongation and differentiation of root cells. The protein phosphatase LIKE SEX FOUR2 (LSF2) has been reported to regulate starch metabolism in *Arabidopsis*, but little is known about the mechanism how LSF2 affect ROS homeostasis. Here, we identified that LSF2 function as a component modulating ROS homeostasis in response to oxidative stress and, thus regulate root development. Compared with wild type *Arabidopsis*, *lsf2-1* mutant exhibited reduced rates of superoxide generation and higher levels of hydrogen peroxide upon oxidative stress treatments. The activities of several antioxidant enzymes, including superoxide dismutase, catalase, and ascorbate peroxidase, were also affected in *lsf2-1* mutant under these oxidative stress conditions. Consequently, *lsf2-1* mutant exhibited the reduced root growth but less inhibition of root hair formation compared to wild type *Arabidopsis* plants. Importantly, protein phosphatase LSF2 interacted with mitogen-activated protein kinase 8 (MPK8), a known component of ROS homeostasis pathways in the cytoplasm. These findings indicated the novel function of LSF2 that controls ROS homeostasis to regulate root development.

Plants absorb water and nutrients from the soil through their roots and root hairs, which sense abiotic and biotic stress during plant life cycle. Reactive oxygen species (ROS) play critical roles in root growth and root hair development in plants. The most common ROS are hydroxyl radical (·OH), superoxide radicals (O₂⁻), and hydrogen peroxide (H₂O₂). Compared with other types of ROS, H₂O₂ is more stable, but O₂⁻ is spontaneously or enzymatically dismutated to H₂O₂. ·OH in plants is directly responsible for cell wall loosening and cell expansion in the root elongation zone. In *Arabidopsis thaliana* roots, O₂⁻ localizes predominantly to the apoplast of cell elongation zone, and H₂O₂ is present mainly in the differentiation zone and cell walls of root hairs. Accumulation of H₂O₂ inhibits root growth but promotes the growth and formation of adventitious roots. Studies in which ROS levels were modified by various treatments have demonstrated that O₂⁻ and H₂O₂ are involved in root growth and root hair development.

Low levels of ROS serve as regulatory signals for the cell, but high levels of ROS, representing byproducts of plant aerobic metabolism produced in response to environmental stresses, are undesirable or toxic. When *Arabidopsis* leaves were exposed to the herbicide paraquat (methyl viologen, MV), H₂O₂ accumulation increased. Salicylic acid (SA) is an oxidative signal inducer that is required for the development of systemic acquired resistance (SAR) against various pathogens through its interaction with NPR3 and NPR4. Plants contain several ROS scavenging enzymes, including superoxide dismutase (SOD), ascorbate peroxidase (APX), and catalase (CAT). The SOD family of metalloenzymes, including iron SOD (Fe-SOD), manganese SOD (Mn-SOD), and copper-zinc...
Oxidative stress affects antioxidant enzyme activities in *lsf2–1* mutant. By determining the levels of ROS in the *lsf2–1* mutant and wild type Col-0 plants, we found that the species of O$_2^•–$ and H$_2$O$_2$ were altered in the *lsf2–1* mutant under external treatment of several oxidative stress inducers. Therefore, we planned to investigate the activities of several enzymes involved in ROS homeostasis such as SOD, APX and CAT in *lsf2–1* mutant under oxidative stress. In plants, SOD represents the first line of defense against excess ROS by catalyzing the dismutation of O$_2^•–$ to H$_2$O$_2$ and molecular oxygen (O$_2$)$_2$. APX is a scavenging peroxidase of H$_2$O$_2$ that uses ascorbate as a reductant and has a high affinity for H$_2$O$_2$. CAT is the key enzyme that scavenges H$_2$O$_2$, with a high reaction rate but low affinity for H$_2$O$_2$. ROS act as either damaging or signaling molecules in plants depending on their levels, which in turn depend on the rates of production and scavenging of ROS.

The precise balance of ROS plays an important role in regulating plant growth and development. Recent studies suggest that ROS homeostasis regulates the transition from cell proliferation to cell differentiation in *Arabidopsis* roots. *Arabidopsis* UPBEAT1 (UPB1) controls the balance between O$_2^•–$ and H$_2$O$_2$ by repressing peroxidases; *upb1* mutant plants contain higher O$_2^•–$ levels and lower H$_2$O$_2$ levels in the root tip compared with the wild type plants, suggesting that ROS homeostasis is altered in these plants, leading to changes in root growth and development. Lack of RHD2/AttrbohC in the *Arabidopsis* rhd2 mutant causes it to have reduced ROS levels and atypical tubulin formation, affecting cell division of the root meristem. PHYTOCHROME AND FLOWERING TIME1 (PFT1), which primarily controls the expression of class III peroxidase genes (such as UPB1) in *Arabidopsis*, regulates the expression of genes encoding redox-active proteins. In the *pft1* mutant, the balance between O$_2^•–$ and H$_2$O$_2$ is altered, resulting in defective root hair differentiation. Over-expressing of PLANT FERREDOXIN-LIKE PROTEIN (PFLP) in *Arabidopsis* increases ROS production and significantly promotes root hair growth. Taken together, these studies support the important role of ROS homeostasis in regulating root growth and development.

In *Arabidopsis*, protein phosphatases play key roles in regulating cellular activities and often affect ROS homeostasis, especially in response to environmental stimuli. There are two known glucan phosphatases in plants: STARCH EXCESS4 (SEX4) and LIKE SEX FOUR2 (LSF2). SEX4 is a major phosphatase that binds to a redox-dependent switch to regulate SEX4 activity. LSF2, which is located in the chloroplast and cytoplasm, scavenges peroxidase of H$_2$O$_2$ that uses ascorbate as a reductant and has a high affinity for H$_2$O$_2$. CAT is the key enzyme that scavenges peroxidase in the SEX4 phosphatase domain that acts as a redox-dependent switch to regulate SEX4 activity. LSF2, which is located in the chloroplast and cytoplasm, specifically hydrolyzes phosphate groups from the C3-position and is involved in transitory starch metabolism. However, no study has proved that LSF2 activity is controlled by redox in plants. In this present study, for the first time, we identify that LSF2 plays a role in ROS homeostasis to regulate root growth and root hair development in *Arabidopsis*.

Results

ROS status is balanced under exogenous oxidative stresses in *lsf2–1* mutant. In order to determine the role of LSF2 in maintenance of ROS levels, we have isolated a homozygous T-DNA inserted mutant *lsf2–1* allele (SAIL line 595_F04 of ABRC; Supplementary Figs. S1a–c) and four independent homozygous *lsf2–1:AtLSF2* complementary transgenic lines (Supplementary Figs. S2a,b). As it is already known that O$_2^•–$ is rapidly converted to H$_2$O$_2$ spontaneously or via dismutation by SOD enzymes. We used XTT as a probe to photometrically measure the generation rate of O$_2^•–$, as XTT is reduced to soluble formazan by O$_2^•–$. Under normal growth conditions, there were no statistical significant differences in O$_2^•–$ generation between *lsf2–1* and Col-0 (Fig. 1a, untreated). When treated with oxidative stress exposure to H$_2$O$_2$, the generation rate of O$_2^•–$ was obviously lower in *lsf2–1* than in Col-0 plants, and these differences were more significant after treatment with exogenous SA or MV (Fig. 1a). We also used NBT as a histochemical probe for O$_2^•–$, as it forms an insoluble blue/violet formazan product in the presence of O$_2^•–$. Strong blue/violet staining (revealing O$_2^•–$) was detected in the elongation zone under abiotic stress treatment, which also extended to the meristematic zone and differentiation zone (Fig. 1b). However, no significant differences were observed between the staining patterns of Col-0, *lsf2–1*, and *lsf2–1:AtLSF2* #1 roots. These results suggest that null mutation of LSF2 in *Arabidopsis* results in more suppressed generation of O$_2^•–$ in response to exogenous oxidative stresses in leaves.

The accumulation of endogenous H$_2$O$_2$ in plants is an early response to environmental stimuli, such as pathogen attack and exogenous application of SA or MV. We measured the contents of endogenous H$_2$O$_2$ using a peroxidase enzyme assay. No significant differences in H$_2$O$_2$ levels were detected between *lsf2–1* and Col-0 plants under normal growth conditions (Fig. 1c, untreated). However, compared with Col-0 plants, *lsf2–1* mutant plants produced higher levels of H$_2$O$_2$ under oxidative stress conditions (Fig. 1c). These results indicate that endogenous H$_2$O$_2$ accumulates much more rapidly in *lsf2–1* than in Col-0 plants under oxidative stress conditions, suggesting that LSF2 is involved in regulating endogenous H$_2$O$_2$ levels exposed to environmental stress. Analysis of LSF2 expression patterns under these exogenous oxidative stresses (SA, H$_2$O$_2$ and MV) in Col-0 plants, the transcriptional expression of LSF2 was strongly induced in leaves but slightly down-regulated in roots (Supplementary Fig. S1d). Therefore, we hypothesized that LSF2 might regulate ROS homeostasis (to a certain extent) by maintaining the balance between O$_2^•–$ levels and H$_2$O$_2$ levels in *Arabidopsis* leaves.
conditions (Fig. 2a,b), suggesting that the activities of Cu/Zn-SODs were more strongly inhibited in the presence of Arabidopsis LSF2 under oxidative stress.

CAT activity is measured by monitoring the extinction of H$_2$O$_2$, whereas APX activity is measured by monitoring ascorbate oxidization. Since endogenous H$_2$O$_2$ levels increased much more strongly in lsf2-1 mutant than in Col-0 plants under oxidative stress conditions, we analyzed the activities of CAT and APX in Col-0, lsf2-1, and lsf2-1:AtLSF2 #1 plants. Under normal growth conditions, no obvious differences in the activities of these enzymes were detected in lsf2-1 mutant compared with Col-0 plants (Fig. 2c,d, untreated). When the seedlings were treated with SA, H$_2$O$_2$, or MV, reduced CAT and APX activity were observed in Col-0, but more significant increases in CAT activity were observed in lsf2-1 (Fig. 2c). However, APX activity was not altered in lsf2-1 in response to the various oxidative stress conditions (Fig. 2d). When seedlings were under oxidative stress, CAT and APX activity were inhibited in the presence of LSF2, whereas CAT activity increased in the absence of this protein. Therefore, CAT, not APX, is mainly responsible for H$_2$O$_2$ scavenging in the absence of Arabidopsis LSF2 under oxidative stress conditions.

Root development is affected by oxidative stresses in lsf2-1 mutant. Based on the altered activities of ROS scavenging enzymes such as SOD, APX and CAT in lsf2-1 and wild type Col-0 plants under several oxidative stress conditions, it seems that LSF2 might be involved in differential regulation of these above mentioned enzymes, which could be the reason for the altered levels of ROS. Previous studies have suggested that ROS level also regulates the root length and root hair development$^{1,24}$. Therefore, we have performed the detail phenotypic assays to investigate the root growth and development under various oxidative stress conditions. Under normal growth conditions, the phenotype of lsf2-1 mutant was not significantly different from that of Col-0. However, when treated with exogenous oxidative stress via exposure to SA, H$_2$O$_2$, or MV, lsf2-1 mutant plants grew poorly compared with Col-0 plants (Fig. 3a,b). Specifically, lsf2-1 roots were shorter than Col-0 roots on medium containing 10μM SA, 0.1 M H$_2$O$_2$, or 2 mM MV (Fig. 3a). The differences in root length between Col-0 and lsf2-1 became more pronounced when grown on medium containing higher concentrations of these agents, such as 50μM SA, 1 mM H$_2$O$_2$, or 20 mM MV (Fig. 3a,b). These findings indicate that the root growth of lsf2-1...
is more strongly inhibited than that of Col-0 under oxidative stress conditions, suggesting that lsf2-1 mutant is hypersensitive to oxidative stress.

No significant differences were found in root hair length or root hair number between lsf2-1 mutant and Col-0 plants under normal growth conditions (Fig. 4a, untreated). However, when treated with oxidative stress via exposure to high concentrations of oxidative stress-inducing agents (50 μM SA, 1 mM H2O2, or 20 nM MV), root hair formation was almost completely inhibited in both Col-0 and lsf2-1 mutant (Fig. 4a). When grown on lower concentrations of these agents (such as 10 μM SA, 0.1 mM H2O2, or 2 nM MV), the root hair density of lsf2-1 was higher than those of Col-0 (Fig. 4b), and the root hairs length of lsf2-1 were much longer than those of Col-0 (Fig. 4c). We also examined the expression of ROOT HAIR DEFECTIVE6 (RHD6), which positively regulates root hair development25. The transcriptional expression of RHD6 was very significantly higher in lsf2-1 than in Col-0 under oxidative stress conditions (Fig. 4d). Together, these results indicate that deficiency of LSF2 in Arabidopsis results in less inhibition of root hair development, as well as less inhibition of RHD6 expression, under oxidative stress conditions compared with normal growth condition (Fig. 4d, untreated).

**Arabidopsis LSF2 interacts with MPK8 in vivo.** Previously, several MAP kinases have been shown to be involved in both biotic and abiotic stresses and affect the ROS homeostasis26. Also dual specificity phosphatases (DSPs) are known to regulate and target several MAP kinases in different abiotic stress condition, such as oxidative stress. To determine if LSF2, a dual specificit protein phosphatase, is also targeting any MAP kinase, we have performed a comprehensive yeast two-hybrid analysis with several MAP kinases (data not shown). Using yeast two-hybrid assays, we found that LSF2 interacted with MPK8 (AT1G18150) (Fig. 5a). To confirm this finding, we purified both full-length LSF2 fused with His, tag and full-length MPK8 fused with glutathione-S-transferase (GST) from E. coli extracts, and then examined the possibility of LSF2 and MPK8 to interact in vitro via a pull down assay and western blotting with anti-GST antibody. The results showed that MPK8 interacted with LSF2 in vitro (Supplementary Fig. S3). To further examine the interaction between LSF2 and MPK8 in vivo, we performed a bimolecular fluorescence complementation (BiFC) assay. Agrobacterium tumefaciens strain EHA105
Figure 3. Phenotypic analysis of root growth in *lsf2-1* mutant and wild type plants under oxidative stress conditions. (a) Seedlings were grown under oxidative stress. (b) The assay of root length of seedlings, which were grown on MS medium containing 50μM SA, 1 mM H₂O₂, or 20 nM MV (n = 20; student's t-test, **P < 0.01).
containing expression binary vectors for nEYFP-LSF2 together with MPK8-cEYFP, or LSF2-cEYFP together with nEYFP-MPK8, were co-infiltrated into N. benthamiana leaves, respectively. Coexpression of nEYFP-LSF2 and MPK8-cEYFP (or LSF2-cEYFP and nEYFP-MPK8) resulted in strong fluorescent signals in the cytoplasm due to reconstitution of the fluorescent marker, EYFP (Fig. 5b). No fluorescent signals were observed when nEYFP-LSF2 or LSF2-cEYFP was coexpressed with cEYFP or nEYFP empty vector, which served as negative controls. Interestingly, the transcriptional expression level of MPK8, not respiratory burst oxidase homolog D (RbohD), was enhanced under these oxidative stress conditions in leaves of lsf2-1 mutant (Fig. 5c; Supplementary Fig. S4).

Discussion
Protein phosphatases (PPs) play important roles in plants. To date, only a few PPs have been characterized in Arabidopsis and other flowering plants. ArRP1 (PPDK regulatory protein) not only exhibits functional properties of protein phosphatases, but it also displays functional properties of protein kinases, as it regulates the activity of pyruvate orthophosphate dikinase (PPDK) via phosphorylation or dephosphorylation of its threonine residue. Interestingly, the plastid-localized enzyme NADP-THIOREDOXIN REDUCTASE C (NTRC) regulates starch synthesis in chloroplasts and amyloplasts in response to either light or sucrose, and it is also involved in maintaining redox homeostasis in plastids of both photosynthetic and nonphotosynthetic tissues, thereby affecting the formation of lateral roots in Arabidopsis. LSF2 is a dual specificity protein phosphatase that displays high activity toward para-nitrophenyl phosphate (pNPP) (Supplementary Fig. S5), and it dephosphorylates glucan to provide access for amylases that release maltose and glucose from starch in Arabidopsis. When the effect of LSF2 null-mutation compared at morphological level, a severity of curly dwarf symptom in lsf2-1 leaves was increased under SA treatment (Supplementary Fig. S6). This suggests the possibility that LSF2 could improve the tolerance to oxidative stresses in plant, which was correlated with the up-regulated expression of LSF2 in wild type leaves under oxidative stress conditions. Hence, the function of LSF2 is not only projected in the root development but...
LSF2 could also be involved in improving plant tolerance to oxidative stresses. The present results suggest that LSF2 also functions in maintaining cellular ROS homeostasis in plants.

Many environmental and genetic factors influence root growth and root hair formation, such as light, calcium, free radicals, and ethylene31. Changes in redox are involved in controlling plant growth and development, and ROS production is associated with root development32. NADPH oxidases and peroxidase have often been shown to be responsible for the production of endogenous \( \text{O}_2^- \) and H\(_2\)O\(_2\) in plants. When H\(_2\)O\(_2\) levels are reduced in plant, increased production of \( \text{O}_2^- \) is an important mechanism that maintains ROS homeostasis11. Redox regulation of AGPase is severely affected in leaves and roots of the Arabidopsis ntrc mutant, and introducing fully functional chloroplasts into this mutant by complementation is both necessary and sufficient for sustaining root growth and lateral root formation, with similar rates to those of the wild type28. The induction of REDOX RESPONSIVE TRANSCRIPTION FACTOR1 (RRTF1) in plants results in ROS accumulation via ROS-generating biotic signals33. Therefore, the balance between \( \text{O}_2^- \) and H\(_2\)O\(_2\) to maintain ROS homeostasis might be involved in root development in plants. Accordingly, in the present study, we investigated the production of \( \text{O}_2^- \) and H\(_2\)O\(_2\) under oxidative stress conditions, the generation rate of \( \text{O}_2^- \) was obviously lower, and the level of endogenous H\(_2\)O\(_2\) was higher, in the lsf2-1 mutant. These changes are consistent with the changes in Cu/Zn-SODs, CAT, and APX activity observed in the lsf2-1 mutant, which led to more inhibition of root growth and less inhibition of root hair development under oxidative stress conditions compared to Col-0 plants. Meanwhile, the expression of some of the marker genes involved in the regulation of root elongation, such as SCN1, CPC and PIN3, were analyzed between Col-0 and lsf2-1 mutant, but no significant difference was found (Supplementary Fig. S7). Furthermore, the expressions of marker genes involved in the regulation of root hairs also were determined by qPCR, including RHD6 gene. RHD6 regulates the development of root hair-forming cells, and Arabidopsis rhd6 mutants have fewer root hairs.

**Figure 5.** Interaction analysis of LSF2 and MPK8 by Y2H and BiFC methodologies. (a) Interaction between LSF2 and MPK8 was assayed by Y2H, C1: \( \text{AD} + \text{BD} \); C2: \( \text{AD-LSF2} + \text{BD} \); C3: \( \text{AD} + \text{BD-MPK8} \); 1: \( \text{AD-LSF2} + \text{BD-MPK8} \). (b) Fluorescence was observed due to complementation of fusion protein LSF2-cEYFP with nEYFP-MPK8, or nEYFP-LSF2 with MPK8-cEYFP, Bars = 20 \( \mu \)m. (c) The relative expression of MPK8 under oxidative stress was detected by qPCR (n = 3; student’s t-test, ***\( P < 0.01 \)).
than the wild type. A recent report showed that ROS production is induced by vanadate treatment and that RHD6 is involved in vanadate-induced root hair initiation. In the present study, the lack of functional LSF2 resulted in less reduced expression of RHD6 under oxidative stress in Arabidopsis, which is consistent with the reduced inhibition of root hair development in the lsf2-1 mutant compared with Col-0 plants. Collectively, the results demonstrate that LSF2 affects root development through modulating ROS homeostasis in Arabidopsis.

Maintaining steady ROS levels appears to require a network involving at least 152 genes in Arabidopsis. PFT1 mainly regulates ROS-related gene expression to control the formation of root hairs via ROS homeostasis. Lower levels of H2O2 are correlated with the reduced expression of POX, and higher levels of O2− are correlated with the increased expression of NADPH. A recent study demonstrated that Mitogen-activated protein kinase kinase kinase 1 (MEKK1) is involved in cellular redox control and integrating ROS homeostasis with plant development and hormone signaling, and MPK4 is the downstream target of MEKK1 in plants. Ca2+ influx and Ca2+/Calmodulins (CaMs) regulate ROS homeostasis and the activities of all antioxidant enzymes. MPK6 mediates Ca2+ influx across the plasma membranes of root cells to control H2O2-induced root elongation in Arabidopsis. Arabidopsis MPK8 interacts with CaM3, CaM4, and CaM7 in the cytoplasm, and is activated by CaMs and mitogen-activated protein kinase kinase 3 (MKK3) to help maintain ROS homeostasis. After mechanical wounding, ROS accumulation increases in the wounding sites of the mpk8 mutant, whereas ROS accumulation decreases in the tissues of MPK8-GFP plants. Furthermore, MPK8 has a conserved phosphorylation motif-TDY and is dual phosphorylated in plants. Furthermore, MPK8 has a conserved phosphorylation motif-TDY and is dual phosphorylated in plants.

Materials and Methods

Plant materials. All plant materials used in this study were in the ecotype Columbia-0 (Col-0) Arabidopsis thaliana background. The lsf2-1 allele is the SAIL line 595_F04 of AT3G10940 and was obtained from the Arabidopsis Biological Resource Center (ABRC). To produce the construct used for functional complementation of the lsf2-1 mutant, the genomic DNA fragment of LSF2 (gLSF2) including the 5′promoter region, coding region, 3′UTR, and following 1282 bp sequence was amplified with the primer pair gLSF2_Fw and gLSF2_Rv (Supplementary Table S1). The resulting fragment was cloned into entry vector pENTR-TOPO (Invitrogen), followed by cloning into binary vector pBA002a. The construct pBA002a-gLSF2 was transformed into the EHA105 strain of Agrobacterium tumefaciens, followed by transformation into lsf2-1 mutant plants by the floral dip method. Transgenic plants harboring gLSF2 were selected on MS medium plates containing 50 μg/ml BASTA. T1 seedlings expressing gLSF2 were grown in soil, and homozygous plants of the four independent complementary lines (lsf2-1:AtLSF2) were identified after three rounds of BASTA application. The insertion site of gLSF2 was determined by genome walking kit (Takara) with three specific primers (SP1, SP2 and SP3; Supplementary Table S1). Arabidopsis seeds were surface-sterilized, and imbibed for 2 days at 4 °C in darkness for stratification, then germinated on MS medium solidified with 0.8% agar in Petri plates. Plants were grown in growth chamber under controlled conditions, 22 °C day/18 °C night with a relative humidity of 60% under long-day conditions (16-h light illumination of 150 μmol·m−2·sec−1, and 8-h dark cycle).

RNA extraction and qPCR analysis. Total RNA was extracted from Arabidopsis using an RNeasy Plant Mini kit (Qiagen). Reverse transcription was performed with 2 μg each of total RNA and oligo (dT) primers using the SuperScript II RT kit (Invitrogen). The cDNA was mixed with SYBR premix Ex Taq (TaKaRa) and gene-specific primers, and amplified in a Bio-Rad CFX96 real-time system. Expression of ACTIN2 was used as a control for cDNA levels. The primer sequences used for qPCR are listed in Supplementary Table S1.

Determination of O2− generation and endogenous H2O2 contents. Pre-treating 2-week-old Col-0, lsf2-1, and lsf2-1:AtLSF2 #1 plants with 2 mM SA, 1 mM H2O2, or 10 μM MV for 30 min, O2− generation of seedlings was measured with the XTT assay, as described by Liszkay, and endogenous H2O2 contents were quantified using a peroxidase enzyme assay according to the method of Youn-II Park. In addition, nitro blue tetrazolium (NBT) staining also was performed as described previously. When NBT reacts with O2−, the color shifts from pale yellow to dark blue, which is used to monitor the generation of superoxide radicals in a histo-chemical staining procedure. The roots were observed under a stereo microscope (Olympus SZX16, Olympus Japan Co., Tokyo, Japan).

Antioxidant enzyme activity assays: superoxide dismutases (SODs), catalase (CAT), and ascorbate peroxidase (APX). Pre-treating 2-week-old Col-0 and lsf2-1 plants with exogenous 2 mM SA for 3 h, 5 mM H2O2 for 30 min, or 30 μM MV for 2 h, SOD activity was detected using NBT staining following electrophoresis as described. Separation of SOD isoforms was performed with each 100 μg protein sample by non-denaturing polyacrylamide gel electrophoresis (PAGE) at 4 °C using a 12% resolving gel and a 5% stacking gel. While, pre-treating 2-week-old Col-0, lsf2-1, and lsf2-1:AtLSF2 #1 plants with exogenous 2 mM SA, 1 mM H2O2, or 10 μM MV for 30 min, CAT activity was determined based on the decrease in absorbance at 240 nm through the extinction of H2O2, and APX activity was determined based on the decrease in absorbance at 290 nm due to ascorbate oxidation.
Root growth and root hair development assays. Col-0, lsf2-1, and lsf2-2:AtLSF2 #1 seeds were surface sterilized and germinated in Petri plates containing MS medium. Exogenous oxidative treatments were performed by adding different concentrations of SA, H₂O₂, or MV (as noted in the figure) to the MS medium. Four- to five-day-old seedlings were transferred individually to five Petri dishes of MS medium. The initial position of the seedling root apex was marked, and the increase in root length was measured after an 8-day incubation. Root growth was also photo-recorded with a digital camera (Nikon D90). The length and density of root hairs were assayed under a stereoscopic microscope (Olympus SZX16). Root growth was calculated with Image J software based on the differences from photographs of the respective time points and plates.

Yeast two-hybrid (Y2H) assay. The Matchmaker GAL4-based two-hybrid system (Clontech) was used to perform yeast two-hybrid assays. The cDNAs encoding full-length LSF2 were cloned into the pGAD424 vector (Clontech) to generate activation domain (AD) constructs. The cDNA encoding full-length MPK8 was cloned into the pGBT9 vector (Clontech) to generate binding domain (BD) constructs. All constructs and empty vector controls were transformed into yeast strain AH109 using the modified lithium acetate method. Co-transformed yeast transformants were selected on plates containing SD/-Leu/-Trp medium and screened on selective medium containing SD/-Leu/-Trp/-His with 20 mM 3-amino-1,2,4-triazole (3-AT) to test protein interactions, and empty vectors pGBT9 and pGAD424 were used as negative controls.

Expression of recombinant proteins and pull-down assay. Full-length LSF2 cDNA was cloned into pET-28a (+) (Novagen) to generate the protein expression vector for biosynthesis of the fusion protein LSF2-His₆. The cDNA encoding full-length MPK8 was inserted into pGEX to generate the translational fusion of GST-MPK8. All recombinant vectors were transformed into E. coli BL21 cells. Isopropyl-β-D-thiogalactoside (IPTG) was used to induce the expression of fusion proteins. GST fusion proteins were purified on glutathione Sepharose™ 4B-Flammmable (GE Health), and His₆ fusion protein was purified on Ni²⁺-nitrilotriacetate (Ni²⁺-NTA) resins (Qiagen). Prey protein LSF2-His₆, and bait fusion protein GST-MPK8 were then added to 1 ml of binding buffer (50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.5% Triton X-100, 0.5 mM β-mercaptoethanol, and 2% proteinase inhibitor cocktail) and incubated with amylase resin beads at 25 °C for 1 h. After incubation, the beads were washed six times with fresh binding buffer. Pull-down proteins were separated by SDS-polyacrylamide gel electrophoresis at 7.5% resolving gel and detected by western blotting using anti-GST antibody (Santa Cruz Biotechnology).

Bimolecular Fluorescence Complementation (BiFC) assay. The vectors pBA-cEYFP(175-end)-N1 (pBAA3032) and pBA-nEYFP (1-174)-C1 (pBAA3036) were used as BiFC Gateway vectors, which respectively generate cDNA-cEYFP and nEYFP-cDNA⁸⁷. The cDNA encoding full-length LSF2 or MPK8 was cloned into the BiFC Gateway vectors, respectively, to test interactions of the proteins in vivo. Recombinant plasmids encoding cEYFP and nEYFP fusions were transformed into competent Agrobacterium tumefaciens (strain EHA105) cells. The EHA105 transformants with recombinant plasmid were suspended in 10 mM MgCl₂ and 150 μM acetosyringone. The mixtures were incubated at 25 °C for at least 3 h without shaking and co-infiltetrated into Nicotiana benthamiana leaves. Microscopic observations of infiltrated plants were performed after 2–3 days. The unfused protein cEYFP or nEYFP was used as a negative control.

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Author Contributions
L.N.S., P.Z., G.-H.L. and Y.-H.Y. designed the experiments. P.Z. performed most of the experiments and drafted the manuscript. L.N.S. performed experiments of in vitro pull down and LSF2 activity. P.Z., L.N.S., C.-Y.T. and Y.Z. performed the Y2H experiment. P.Z. and J.Y. performed BiFC experiments. P.Z., G.-H.L., G.K.P. and Y.-H.Y. analyzed the data. G.K.P., G.-H.L., J.S., W.L., Z.H. and J.Q. revised the manuscript.

Additional Information
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