Zinc Transporter ZmLAZ1-4 Modulates Zinc Homeostasis on Plasma and Vacuolar Membrane in Maize

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Zinc is an essential micronutrient for plant growth and development, and functions as a cofactor for hundreds of transcription factors and enzymes in numerous biological processes. Zinc deficiency is common abiotic stress resulting in yield loss and quality deterioration of crops, but zinc excess causes toxicity for biological systems. In plants, zinc homeostasis is tightly modulated by zinc transporters and binding compounds that uptake/release, transport, localize, and store zinc, as well as their upstream regulators. Lazarus 1 (LAZ1), a member of DUF300 protein family, functions as transmembrane organic solute transporter in vertebrates. However, the function of LAZ1 in plants is still obscure. In the present study, the ZmLAZ1-4 protein was confirmed to bind to zinc ions by bioinformatic prediction and thermal shift assay. Heterologous expression of ZmLAZ1-4 in the zinc-sensitive yeast mutant, Arabidopsis, and maize significantly facilitated the accumulation of Zn$^{2+}$ in transgenic lines, respectively. The result of subcellular localization exhibited that ZmLAZ1-4 was localized on the plasma and vacuolar membrane, as well as chloroplast. Moreover, the ZmLAZ1-4 gene was negatively co-expressed with ZmBES1/BZR1-11 gene through co-expression and real-time quantitative PCR analysis. The results of yeast one-hybrid and dual-luciferase assay suggested that ZmBES1/BZR1-11 could bind to ZmLAZ1-4 promoter to inhibit its transcription. All results indicated that ZmLAZ1-4 was a novel zinc transporter on plasma and vacuolar membrane, and transported zinc under negative regulation of the ZmBES1/BZR1-11 transcription factor. The study provides insights into further underlying the mechanism of ZmLAZ1-4 regulating zinc homeostasis.

Keywords: maize, ZmLAZ1-4, zinc transport, tonoplast, transcriptional regulation

INTRODUCTION

Zinc (Zn) is an essential micronutrient for plant growth and development, and functions as a cofactor for hundreds of transcription factors and enzymes in numerous biological processes, such as chlorophyll biosynthesis, gene expression, signal transduction, and stress response (Palmer and Guerinot, 2009; Zlobin, 2021). Zn deficiency is common abiotic stress resulting in production
loss and quality deterioration of crops, but Zn excess causes toxicity for biological systems (Grotz et al., 1998; Arrivault et al., 2006; Palmer and Guerinot, 2009). However, Zn deficiency is far more frequent than toxicity, because Zn content in the soils has low availability for plants (Alvarez and Rico, 2003; Cakmak, 2008). Zn toxicity only occurs on polluted soils containing excessive Zn in mining or industrial areas (Mossa et al., 2020). In plants, the excessive Zn is usually stored in the vacuole to avoid toxicity (Martinova et al., 2007). Usually, Zn homeostasis is tightly modulated by Zn transporters and binding compounds that uptake/release, transport, localize, and store Zn within the whole plant as well as within individual tissues, cells, and cellular compartments (Palmer and Guerinot, 2009; Zlobin, 2021). Some metal tolerance proteins (MTPs) and heavy metal ATPases (HMAs) localize on the vacuolar membrane and modulate Zn homeostasis as a Zn sensor and transporter (Arrivault et al., 2006; Menguer et al., 2013). In Arabidopsis, two LAZ1 proteins are found to maintain vacuole integrity and mediate brassinosteroid (BR) signaling and localized on the vacuolar membrane (Liu et al., 2018). In our previous study, we cloned eight members of the ZmLAZ1 gene family from maize and found their differential expression among different organs, developmental stages, and under abiotic stresses, implying their functional diversity (Liu et al., 2020). In the present study, we demonstrated that the ZmLAZ1-4 protein was distinct from the other seven members, and it functioned as a zinc transporter on plasma and vacuolar membrane and modulated zinc homeostasis under the negative regulation of BR signaling transcription factor ZmBES1/BZR1-11.

**MATERIALS AND METHODS**

**Substrate Prediction and Thermal Shift Assay**

To predict candidate substrates of ZmLAZ1 proteins, the amino acid sequences of eight ZmLAZ1 members were submitted to the SWISS-MODEL software1 to get a protein model structure file in PDB format. Then the PDB file of each ZmLAZ1 was searched against the RCSB PDB software2 to get putative substrates. The coding sequences (CDSs) of the ZmLAZ1-4 and ZmLAZ1-8 genes were amplified from pMD19-T-ZmLAZ1-4 and pMD19-T-ZmLAZ1-8 (Liu et al., 2020) by PCR primers CDS1-4F/CDS1-4R and CDS1-8F/CDS1-8R (Supplementary Table 1), respectively, and inserted into His-tagged prokaryotic expression vector pET-32a (Takara, Osaka, Japan) by using ClonExpress II One Step Cloning Kit (Vazyme Biotech, Nanjing, China). The construct was introduced into Escherichia coli BL21 (DE3), screened on Luria-Bertani (LB) plates containing 100 mg/ml ampicillin, and grown in LB medium at 37°C overnight, purified by using Ni-TED 1 ml Sefinose (TM) Column (Sangon Biotech, Shanghai, China), detected by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), quantified in NanoDrop One Microvolume UV–Vis Spectrophotometer (Thermo Fisher Scientific, Waltham, MA, United States), and diluted to 1 µg/µl with 10% dimethyl sulfoxide (DMSO).

As described by Huynh and Partch (2015) with minor modification, 2 µl of 10% DMSO, 6 µg of the purified protein, 2 µl of 10 × SYPRO orange, and 0 (blank control) and 200 µM

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1. https://swissmodel.expasy.org/
2. https://www.rcsb.org/
of each predicted substrate were added into each of three wells of a 96-well PCR plate. In CFX Connect Real-Time PCR Detection System (Bio-Rad, Hercules, CA, United States), the sampled plate was equilibrated at 25°C for 5 min and then ramped up to a final temperature of 95°C in increments of 1°C. Fluorescence was read every 0.2°C ramping up. The change rates of relative fluorescence units (RFUs) with time (T) [-d(RFU)/dT] were plotted vs. the temperature to generate melting curves of ZmLA1-4 incubated with each predicted substrate.

**Zinc Transport Assay in Zinc-Sensitive Yeast Mutant**

The CDS of ZmLAZ1-4 was amplified from pMD19-T-ZmLAZ1-4 plasmid (Liu et al., 2020) by using primers pYES2F/pYES2R (Supplementary Table 1) and used for construction of yeast expression vector pYES2-ZmLAZ1-4 by using ClonExpress® II One Step Cloning Kit (Vazyme Biotech, Nanjing, China). The pYES2-ZmLAZ1-4 and empty vector pYES2 were transformed into yeast Zn-sensitive mutant Δzrc1 (Miyabe et al., 2001) and wild-type (WT) strain BY4743, respectively. The positively transformed lines were selected on synthetic dropout medium (SD) plates containing 2% galactose (w/v) and without uracil (Ura), identified by PCR amplification using primers Non-Term1-4F/Non-Term1-4R and AtTIP2F/AtTIP2R (Loque et al., 2005) was amplified from pMD19-T-ZmLAZ1-4 plasmid (Liu et al., 2020) and Arabidopsis cDNA using primers Non-Term1-4F/Non-Term1-4R and AtTIP2F/AtTIP2R (Supplementary Table 1) and used for construction of transient expression vector 35S-ZmLAZ1-4-eGFP and 35S-mCherry-AtTIP2 using ClonExpress® II One Step Cloning Kit (Vazyme Biotech, Nanjing, China), respectively. Subsequently, the 35S-ZmLAZ1-4-eGFP, 35S-mCherry-AtTIP2 of tonoplast maker, 35S-mCherry-OsRac3 of plasma membrane marker (donated by professor Shuangcheng Li, Tao et al., 2021), and empty vector 35S-eGFP (blank control) were introduced into Agrobacterium tumefaciens strain GV3101, respectively.

Maize mesophyll protoplasts were prepared with etiolated leaves and co-transfected with 35S-ZmLAZ1-4-eGFP and 35S-mCherry-OsRac3 as well as 35S-eGFP and 35S-mCherry-OsRac3

**Subcellular Localization**

The transmembrane domains of LAZ1-4 were predicted by the TMHMM v. 2.0 software.3 The CDS without termination codons of ZmLAZ1-4 and tonoplast maker gene AtTIP2 (Loque et al., 2005) was amplified from pMD19-T-ZmLAZ1-4 plasmid (Liu et al., 2020) and Arabidopsis cDNA using primers Term1-4F/Non-Term1-4R and AtTIP2F/AtTIP2R (Supplementary Table 1) and used for construction of transient expression vector 35S-ZmLAZ1-4-eGFP and 35S-mCherry-AtTIP2 using ClonExpress® II One Step Cloning Kit (Vazyme Biotech, Nanjing, China), respectively. Subsequently, the 35S-ZmLAZ1-4-eGFP, 35S-mCherry-AtTIP2 of tonoplast maker, 35S-mCherry-OsRac3 of plasma membrane marker (donated by professor Shuangcheng Li, Tao et al., 2021), and empty vector 35S-eGFP (blank control) were introduced into Agrobacterium tumefaciens strain GV3101, respectively.

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**Transformation and Phenotyping of Arabidopsis and Maize**

Overexpression vector (pC2300-35S-ZmLAZ1-4) of ZmLAZ1-4 was constructed as above and introduced into Agrobacterium tumefaciens strain GV3101. Positive strains were identified and used for transformation of WT Arabidopsis thaliana by floral dip. Transgenic lines were selected on kanamycin 1/2 MS plates and identified by PCR with primers LAZ4F1/LAZ4R1 (Supplementary Table 1). Referring to Kawachi et al. (2009), each homozygous line was grouped into three replicates and grown on 1/2 MS zinc deficiency plates (control) with 5 and 50 µM ZnSO₄ at 22°C temperature, 50% humidity, and 16 h light of 120 µE m⁻² s⁻¹ illumination intensity/8 h dark period for 2 weeks. After photographing for the phenotype, the seedlings were dried at 60°C for 48 h, weighed for biomass, and digested in 80% nitric acid at 250°C overnight. The digested solution was diluted with ddH₂O and used for measurement of Zn²⁺ content by Inductively Coupled Plasma-Mass Spectrometry (Thermo Fisher Scientific, Waltham, MA, United States).

Overexpression vector (pZZ00026-Ubi-ZmLAZ1-4-Tnos) of ZmLAZ1-4 was constructed as above and used to transform embryonic calli isolated from maize inbred line B73 by Agrobacterium mediation. Positive calli were screened on H6 medium with 0.06% (v/v, effective concentration) Basta herbicide. Regenerated plantlets were screened by PAT/bar EPSPS LFD Strip kit (Youlong, Shanghai, China) according to the manufacturer’s instruction and identified by PCR with primers LAZ4F2/LAZ4R2 (Supplementary Table 1) and real-time quantitative PCR (RT-qPCR) with primers LAZ4F3/LAZ4R3 (Supplementary Table 1). Homozygous T₃ lines and WT were grown in vermiculite with Zn dropout Hoagland’s nutrient solution (Coolaber, Beijing, China) at 28°C and 300 µm.ₑ⁻⁻² s⁻¹ illumination intensity for 16 h/20°C dark for 8 h. Referring to Kawachi et al. (2009), at three-leaf stage, the seedlings of each line were grouped into three replicates, treated with 5 and 50 µM ZnSO₄ for 3 weeks, then photographed and dried at 60°C for 72 h, and weighed for biomass and used to measure Zn²⁺ content as above.

**Subcellular Localization**

The transmembrane domains of LAZ1-4 were predicted by the TMHMM v. 2.0 software.3 The CDS without termination codons of ZmLAZ1-4 and tonoplast maker gene AtTIP2 (Loque et al., 2005) was amplified from pMD19-T-ZmLAZ1-4 plasmid (Liu et al., 2020) and Arabidopsis cDNA using primers Term1-4F/Non-Term1-4R and AtTIP2F/AtTIP2R (Supplementary Table 1) and used for construction of transient expression vector 35S-ZmLAZ1-4-eGFP and 35S-mCherry-AtTIP2 using ClonExpress® II One Step Cloning Kit (Vazyme Biotech, Nanjing, China), respectively. Subsequently, the 35S-ZmLAZ1-4-eGFP, 35S-mCherry-AtTIP2 of tonoplast maker, 35S-mCherry-OsRac3 of plasma membrane marker (donated by professor Shuangcheng Li, Tao et al., 2021), and empty vector 35S-eGFP (blank control) were introduced into Agrobacterium tumefaciens strain GV3101, respectively.

Maize mesophyll protoplasts were prepared with etiolated leaves and co-transfected with 35S-ZmLAZ1-4-eGFP and 35S-mCherry-OsRac3

**TABLE 1 | Predicted substrates for the ZmLAZ1 family.**

| Protein | Substrate |
|---------|-----------|
| ZmLAZ1-1 | – |
| ZmLAZ1-2 | MAN | BMA | NAG | CE |
| ZmLAZ1-3 | – |
| ZmLAZ1-4 | Zn²⁺ | MAN | BMA | Mg²⁺ | NAG | OPA |
| ZmLAZ1-5 | MAN | BMA | NAG | CE |
| ZmLAZ1-7 | Dodecane | Retinal | Decane | PPC |
| ZmLAZ1-8 | ESA | Zn²⁺ | Enolate | Ca²⁺ | TPP + Fe | OBG |
| ZmLAZ1-9 | – |

MAN, α-D-mannose; BMA, β-D-mannose; NAG, N-acetyl-D-glucosamine; CE, Cholesterol; PPC, Phosphocholine; ESA, ethanolsulfonic acid; OPA, Oxyphosphonic acid; TPP, toporphyrin; OBG, octyl-β-octylglycoside—means no predicted substrate.

3http://www.cbs.dtu.dk/services/TMHMM
FIGURE 1 | Thermal shift assay of ZmLA1-4 incubated with predicted substrates.

as blank control. After incubation at 25°C in dark for 12 h, the protoplasts were used to observe the fluorescence of eGFP and OsRac3 under laser scanning confocal microscope LSM 800 with 488 and 584 nm laser channel (Zeiss, Oberkochen, Germany), respectively.

The combination of 35S-ZmLAZ1-4-eGFP and 35S-mCherry-OsRac3, 35S-ZmLAZ1-4-eGFP and 35S-mCherry-AtTIP2, as well as 35S-eGFP and 35S-mCherry-OsRac3, and 35S-eGFP and 35S-mCherry-AtTIP2 plasmid was mixed with 0.1 M spermidine and 2.5 M CaCl₂ and precipitated onto gold particles (ϕ = 60 µm), respectively. Onion bulbs were surface sterilized with 75% ethanol. The fifth scales without pigment were cut into 2 cm × 2 cm, incubated on MS medium for 4 h, and bombarded in helium biolistic gun (Bio-Rad, Hercules, CA, United States) with above gold particles. After filtration at 28°C under dark for 24 h, the bombarded onion scales were used to observe the fluorescence of eGFP, OsRac3, and AtTIP2 under the same microscope.

The Agrobacterium strains harboring 35S-ZmLAZ1-4-eGFP and 35S-eGFP were infiltrated into the abaxial leaf surface of 3-week-old plants of Nicotiana benthamiana, respectively. After incubation at 22°C and 14 light/10 dark for 24 h, the infiltrated leaves were used to observe the fluorescence of eGFP and autofluorescence of chloroplasts under the same microscope.

Co-expression and Real-Time Quantitative PCR Analysis
The transcriptomic data of maize inbred line B73 were downloaded from MaizeGDB⁴ and used for co-expression analysis with ZmLAZ1-4 by using a Perl script (Supplementary Data Set 1). The correlation coefficient was set as > 0.9 and < −0.9. Among the candidates, only the ZmBES1/BZR1-11 gene encoded transcription factor, co-expressed with ZmLAZ1-4 (correlation coefficient is −0.93), and used for RT-qPCR analysis. The seeds of inbred line B73 were surface sterilized with 30% H₂O₂, germinated in petri dish, and transplanted into a plastic mesh grid for hydroponic culture at 28°C under a photoperiod of 14 h light/10 h dark. At the three-leaf stage, the seedlings were subjected to the treatment of 5 µM ZnSO₄. At the 0 (control), 1st, 2nd, and 3rd day of treatment, the whole plant was sampled, ground in liquid nitrogen, and used for total RNA extraction by using RNAiso plus kit (TaKaRa, Osaka, Japan). After removing probable genomic DNA contamination by using RNase-free DNase (TaKaRa, Osaka, Japan), these samples were quantified on NanoDrop 2000 (Thermo Fisher Scientific, Waltham, MA, United States) and reverse transcribed into cDNA by using PrimeScript™ reagent kit (TaKaRa, Osaka, Japan). The RT-qPCR was performed as described by Sun et al. (2020). The ZmGAPDH was used as reference. The primers are listed in Supplementary Table 1.

Yeast One Hybrid and Dual Luciferase Assay
The CDS of ZmBES1/BZR1-11 was amplified with primers pGADT7F/pGADT7R (Supplementary Table 1) and used to construct vector pGADT7-ZmBES1/BZR1-11. The cis-acting elements bound by ZmBES1/BZR1-11 in ZmLAZ1-4 promoter were predicted by PlantCARE.⁵ The sequence (−1 to

⁴https://www.maizegdb.org

⁵http://bioinformatics.psb.ugent.be/webtools/plantcare/html/
The promoter sequence (−1 to −1,100 bp) of ZmLAZ1-4 promoter (pZmLAZ1-4) was amplified with specific primers pGreenIIF/pGreenIIR (Supplementary Table 1) and inserted into pGreenI-0800-LUC plasmid to drive firefly luciferase gene (LUC) and generate reporter vector pZmLAZ1-4-LUC. The Renilla luciferase gene REN driven by 35S promoter in pZmLAZ1-4-LUC plasmid was used as internal reference. The CDS of the ZmBESI/BZR1-11 gene was amplified with primers pCAMBIA2300F/pCAMBIA2300R (Supplementary Table 1) and inserted into pCAMBIA2300-35S-eGFP plasmid to create effector vector 35S-ZmBESI/BZR1-11. The reporter and the effector vectors were introduced into Agrobacterium strain GV3101, respectively, and used for co-infiltration of Nicotiana benthamiana leaves. After incubated at 22°C and 14 light/10 dark for 3 days, the leaves were visualized for LUC signal in ChemiDoc™ Imaging System (Bio-Rad, Hercules, CA, United States). The relative activities of LUC and REN were determined in a dual-luciferase reporter assay system (Thermo...
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**FIGURE 3** | Phenotype and Zn$^{2+}$ content of transgenic Arabidopsis. (A) Phenotype. (B) Zn$^{2+}$ content. The seeds of every line were sterilized and grown on 1/2 MS zinc deficiency plates (control) with 5 and 50 µM ZnSO$_4$ for 2 weeks, respectively. WT, wild type. ZmLA1-4 I and ZmLA1-4 II represent homozygous lines. *p < 0.05.

Fisher Scientific, Waltham, MA, United States) and used to calculate relative LUC activity (LUC/REN).

**Statistical Analysis**

All experiments were performed with three replicates. The data were shown as mean ± standard deviation and analyzed using Student's t-test at *p < 0.05 and **p < 0.01 level.

**RESULTS**

**ZmLAZ1-4 Specifically Binds to Zinc**

By the RCSBPDB software as described by Wang et al. (2001), substrates of eight ZmLAZ1 members were mainly predicted to be organic solutes including α-D-mannose, β-D-mannose, N-acetyl-D-glucosamine, cholesterol, phosphocholine, ethanesulfonic acid, phosphinic acid, toporphyrin, and octyl-β-octylglucoside. However, only ZmLAZ1-4 (Zm00001d012921) and ZmLAZ1-8 (Zm00001d036361) were predicted to combine inorganic ions containing zinc (Zn$^{2+}$), magnesium (Mg$^{2+}$), and calcium (Ca$^{2+}$) (Table 1). During many times of prokaryotic expression, the ZmLAZ1-8 protein was not successfully purified (Supplementary Figure 1). Therefore, the ZmLAZ1-4 was used for further study.

In the thermal shift assay, melting temperature (Tm) of the ZmLAZ1-4 protein incubated with ZnCl$_2$ and ZnSO$_4$ was 4.7 and 4.5°C lower than that of ZmLAZ1-4 incubated alone (blank control), respectively, whereas Tm of ZmLAZ1-4 incubated with other predicted substrates kept same value with blank control (Figure 1). This result suggested that zinc ion was candidate substrate of ZmLAZ1-4.

**ZmLAZ1-4 Transports Zinc in Yeast, Arabidopsis, and Maize**

Under non-stress (0 mM Zn$^{2+}$), the diluted colonies and growth curves of OD$_{600}$ showed no significant difference among Δzrc1 mutant transformed by the ZmLAZ1-4 gene, and Δzrc1 and WT transformed by empty vector pYES2. Under 2 mM Zn$^{2+}$ stress, the difference was significant among these three lines (Figures 2A,B). The complementation of ZmLAZ1-4 significantly inhibited the growth of Zn-sensitive mutant Δzrc1. The Zn$^{2+}$ concentration of Δzrc1 transformed by ZmLAZ1-4 was significantly higher than that of Δzrc1 and WT transformed by empty vector pYES2 (Figure 2C), suggesting that ZmLAZ1-4 could transport Zn$^{2+}$ into cells.

Two homozygous T$_3$ Arabidopsis lines overexpressing ZmLAZ1-4 were screened on kanamycin 1/2 MS plates and identified by PCR amplification (Supplementary Figure 1). Under 0, 5, and 50 µM ZnSO$_4$ treatments, the growth phenotype of T$_3$ lines showed no obvious difference compared to WT (Figure 3A). However, Zn$^{2+}$ content of transgenic lines was significantly higher than WT under 5 and 50 µM ZnSO$_4$
FIGURE 4 | Phenotype of transgenic maize under ZnSO$_4$ treatment. (A) Phenotype. (B) Biomass. (C) Zn$^{2+}$ content. At three-leaf stage, the seedlings of each line were grouped into three replicates, treated with 5 and 50 µM ZnSO$_4$ for 3 weeks, then photographed and dried at 60°C for 72 h, and weighed for biomass and used to measure Zn$^{2+}$ content. The biomass of three seedlings of every line was shown. B73, the untransformed control. Line 4, 6, 7, and 10 are homozygous T$_3$ lines. *p < 0.05.

treatments, while only trace content was measured under 0 µM ZnSO$_4$ treatment (Figure 3B).

By *A. tumefaciens*-mediated embryonic calli transformation, from the positive transgenic calli harboring *ZmLAZ1-4*, ten plantlets were regenerated and four homozygous T$_3$ maize lines overexpressing *ZmLAZ1-4* were identified by PAT/bar EPSPS LFD Strips (Supplementary Figure 3), PCR amplification (Supplementary Figure 4), and RT-qPCR (Supplementary Figure 5). Four homozygous lines and WT were grown in vermiculite with Zn deficient Hoagland’s nutrient solution. After 3 weeks of 5 and 50 µM ZnSO$_4$ treatments, the growth phenotype and biomass of all transgenic lines showed different compared with WT (Figures 4A,B). However, the Zn$^{2+}$ content of all transgenic lines was significantly higher than WT under 5 and 50 µM ZnSO$_4$ treatments (Figure 4C). The above results indicated that *ZmLAZ1-4* functioned as a Zn transporter.

**ZmLAZ1-4 Localized on Plasma and Vacuolar Membranes**

By the TMHMM software, seven transmembrane domains were predicted during *ZmLAZ1-4* protein (Supplementary Figure 6). As shown in Figure 5, the GFP fluorescence was observed in the cytoplasm and nucleus in maize protoplasts, and onion
FIGURE 5 | Subcellular localization of ZmLAZ1-4 in maize protoplasts (A) and onion cells (B). Scale bar is 50 µm.
cells transfected by empty vector 35S-eGFP. However, the GFP fluorescence from the fusion protein (35S-ZmLAZ1-4-GFP) was merged with red fluorescence of the plasma membrane marker OsRac3, tonoplast marker AtTIP2, and autofluorescence of chloroplasts. Especially when AtTIP2 was used as a maker, it could be clearly seen that ZmLAZ-4 was localized on the tonoplast. Furthermore, ZmLAZ1-4 was also localized to chloroplast (Supplementary Figure 7). These results indicated the subcellular localization of the ZmLAZ1-4 protein on the plasma and vacuolar membrane.

ZmLAZ1-4 Is Negatively Regulated by ZmBES1/BZR1-11

In order to explore the mechanism of ZmLAZ1-4 regulating Zn transport, the co-expression analysis was conducted. The results showed that there were 27 genes co-expressed with ZmLAZ1-4 with correlation coefficient $> 0.9$ or $< -0.9$, and only ZmBES1/BZR1-11 among these candidates encoded transcription factor and negatively co-expressed with ZmLAZ1-4 (Supplementary Table 2). The result of RT-qPCR likewise showed that the expression of ZmLAZ1-4 and ZmBES1/BZR1-11 was significantly downregulated and upregulated by Zn deficiency ($5 \mu$M), respectively (Figure 6). It was predicted that there were six E-boxes (CAXXTG) of BES1/BZR1 binding element (Yin et al., 2005) during ZmLAZ1-4 promoter by PlantCARE. Hence, the yeast one-hybrid (Y1H) was performed. As shown in Figure 7, on Leu dropout SD medium, the growth of Y1H gold strain co-transformed by empty prey vector pGADT7 and pAbAi-pZmLAZ1-4 harboring six E-boxes elements of ZmLAZ1-4 promoter was inhibited by 200 ng/ml AbA, whereas the Y1H gold strain co-transformed by pGADT7-ZmBES1/BZR1-11 and pAbAi-pZmLAZ1-4 formed few colonies, indicating that the ZmBES1/BZR1-11 transcription factor could bind to ZmLAZ1-4 promoter. The result was further verified by dual-luciferase assay in vivo. The relative LUC activity (LUC/REN) of leaves co-infiltrated by reporter vector ZmLAZ1-4-LUC and effector vector 35S-ZmBES1/BZR1-11 was significantly lower than that of control (Figure 7B). These results indicate that the ZmBES1/BZR1-11 transcription factor binds to ZmLAZ1-4 promoter to inhibit ZmLAZ1-4 transcription.

DISCUSSION

The eight ZmLAZ1 members were grouped into a family in phylogenetic analysis because of their sequence similarity, especially their conserved DUF300 domain (Malinovsky et al., 2010; Liu et al., 2020). In the present study, only ZmLAZ1-4 and ZmLAZ1-8 were predicted to combine metal ions including Zn$^{2+}$, Mg$^{2+}$, or Ca$^{2+}$ (Table 1). During prokaryotic expression, ZmLAZ1-8 was not successfully purified (Supplementary Figure 1). Eukaryotic membrane proteins are often difficult to be purified (Newstead et al., 2007). Therefore, the combination of ZmLAZ1-4 to predicted substrates was verified by thermal shift assay (Figure 1). Even in ZIP family, only some members were identified as Zn transporters (Grotz et al., 1998; Ishimaru et al., 2007; Evens et al., 2017; Yang et al., 2020). The other members might function as transporters of other divalent ions (Miller et al., 2013). The overexpression of ZmLAZ1-4 in yeast mutant, Arabidopsis, and maize significantly increased Zn uptake (Figures 2–4), suggesting that the ZmLAZ1-4 protein was involved in Zn uptake in maize.

In our previous study, ZmLAZ1-4 was predicted to localize on chloroplast, plasmalemma, cytoplasm, and endoplasmic reticulum (Liu et al., 2020). The subcellular localization showed that ZmLAZ1-4 functioned on plasma and vacuolar membrane, as well as chloroplast using tonoplast maker AtTIP2 and plasma membrane marker OsRac3 (Figure 5 and Supplementary Figure 7), which well confirmed the tonoplast and plasma membrane localization (Loque et al., 2005; Tao et al., 2021). Hence, they could be used as a marker in our study. The phenomenon was similar with Mg$^{2+}$ transporter AtMRS2 showing different intracellular localization patterns in yeast and chloroplast localization, and PI transporter PHT2;1 localizing to mitochondria, plasma membrane, endoplasmic reticulum, and chloroplast (Wayne and Maria, 2002; Drummond et al., 2006). In Arabidopsis, two LAZ1 proteins were also localized on plasma and vacuolar membrane, but no specific marker was used for chloroplast localization (Malinovsky et al., 2010). Our result suggests that ZmLAZ1-4 functions on the plasma membrane and uptakes Zn from the soil, and transports Zn into vacuole. But the mechanism of ZmLAZ1-4 acting on chloroplast remains unclear. Before this study, Zn transport across chloroplast and vacuolar membrane was well documented to be mediated by HMA, MTP, and Oryza sativa Zn transporter (OTZ) proteins (Kobae et al., 2004; Arrivault et al., 2006; Martinosta et al., 2007; Moreno et al., 2008; Kim et al., 2009; Morel et al., 2009; Lan et al., 2013; Mengue et al., 2013; Tanaka et al., 2015). Some endoplasmic reticulum-localized and Golgi apparatus-localized zinc transporters were also involved in Zn homeostasis by controlling the release of zinc into cytosol (Fujivara et al., 2015; Adulcikas et al., 2018; Wang et al., 2021). Our result of subcellular localization could not rule out the possibility of endoplasmic reticulum and Golgi apparatus localization of ZmLAZ1-4 (Supplementary Figure 7). This will be explored in further study.

Among the 27 genes co-expressing with ZmLAZ1-4, only ZmBES1/BZR1-11 encoded transcription factor and negatively co-expressed with ZmLAZ1-4 (Figure 6 and Supplementary Table 2). The ZmBES1/BZR1-11 can bind
FIGURE 7 | Confirmation of ZmBES1/BZR1-11 binding to ZmLAZ1-4 promoter. The Y1H (A) and dual-luciferase assay (B) were performed to verify the binding and negative regulation of ZmLAZ1-4 by ZmBES1/BZR1-11 transcription factor. The tobacco leaves were co-infiltrated by 35S-ZmBES1/BZR1-11 and pZmLAZ1-4-LUC, incubated at 22°C and 14 light/10 dark for 3 days, visualized for LUC signal, and used to measure activity LUC and REN. **p < 0.01.

FIGURE 8 | Signaling diagram of zinc homeostasis in maize. The black words, lines, and arrows indicate signaling pathways previously reported. The red words, lines, and arrows show signaling pathway demonstrated in the study.

to E-boxes (CAXXTG) element in ZmLAZ1-4 promoter to inhibit ZmLAZ1-4 transcription (Yin et al., 2005), which was verified by Y1H and dual-luciferase assay (Figure 7). But previous studies exhibited that BES1/BZR1 transcription factor responds to BR induction and regulates the expression of BR-responsive genes (Yin et al., 2005; Yu et al., 2018), and Arabidopsis LAZ1 proteins localized on plasma and vacuolar membrane also mediated BR signaling (Malinovsky et al., 2010). It could be concluded that the ZmLAZ1-4 protein functioned as a Zn$^{2+}$ transporter on plasma and vacuolar membrane, and chloroplast to modulate Zn homeostasis in maize. The expression of ZmLAZ1-4 was negatively regulated by ZmBES1/BZR1-11 transcription factor. The results of this study indicated that ZmLAZ1-4 was a novel zinc transporter distinct from the previously documented Zn transporters ZIP, ZRT, IRT, NRAMP, etc. (Grotz et al., 1998; Ishimaru et al., 2007;
Milner et al., 2013; Evens et al., 2017; Yang et al., 2020), as plotted in a signaling diagram of zinc homeostasis together with the previously reported evidence (Figure 8).

CONCLUSION

The ZmLAZ1-4 protein is a novel zinc transporter that transports zinc ions across plasma and vacuolar membrane and modulates zinc homeostasis under the negative regulation of ZmBES1/BZR1-11 transcription factor.

DATA AVAILABILITY STATEMENT

Publicly available datasets were analyzed in this study. These data can be found here: wap.maizegdb.org, Zea_mays.AGPv4.32gff3.

AUTHOR CONTRIBUTIONS

FF and WL conceived and supervised the research. BL, HY, QyY, LD, FS, JQ, WF, and QqY performed the experiments. BL and WL drafted the manuscript. BL and HY revised the manuscript. All authors interpreted and discussed the data.

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SUPPLEMENTARY MATERIAL

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Figure 8
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