An endoplasmic reticulum–localized cytochrome b<sub>5</sub> regulates high-affinity K<sup>+</sup> transport in response to salt stress in rice

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Potassium (K<sup>+</sup>) is an essential element for growth and development in both animals and plants, while high levels of environmental sodium (Na<sup>+</sup>) represent a threat to most plants. The uptake of K<sup>+</sup> from high-saline environments is an essential mechanism to maintain intracellular K<sup>+</sup>/Na<sup>+</sup> homeostasis, which can help reduce toxicity caused by Na<sup>+</sup> accumulation, thereby improving the salt tolerance of plants. However, the mechanisms and regulation of K<sup>+</sup>-uptake during salt stress remain poorly understood. In this study, we identified an endoplasmic reticulum–localized cytochrome b<sub>5</sub> (OsCYB5-2) that interacted with a high-affinity K<sup>+</sup> transporter (OsHAK21) at the plasma membrane. The association of OsCYB5-2 with the OsHAK21 transporter caused an increase in transporter activity by enhancing the apparent affinity for K<sup>+</sup>-binding but not Na<sup>+</sup>-binding. Heme binding to OsCYB5-2 was essential for the regulation of OsHAK21. High salinity directly triggered the OsHAK21–OsCYB5-2 interaction, promoting OsHAK21-mediated K<sup>+</sup>-uptake and restricting Na<sup>+</sup> entry into cells; this maintained intracellular K<sup>+</sup>/Na<sup>+</sup> homeostasis in rice cells. Finally, overexpression of OsCYB5-2 increased OsHAK21-mediated K<sup>+</sup> transport and improved salt tolerance in rice seedlings. This study revealed a posttranslational regulatory mechanism for HAK transporter activity mediated by a cytochrome b<sub>5</sub> and highlighted the coordinated action of two proteins to perceive Na<sup>+</sup> in response to salt stress.

**Significance**

High-affinity K<sup>+</sup> (HAK) transporter-mediated K<sup>+</sup>-uptake has an important role when plants are subjected to stresses. This work identifies a mechanism of HAK regulation. The affinity of HAK at the plasma membrane for K<sup>+</sup> depends on the binding of a cytochrome (CYB5) protein at the endoplasmic reticulum. This improves K<sup>+</sup>-uptake and the ability of plants to survive under saline conditions. The HAK–CYB5 interaction not only constitutes a mechanism of HAK regulation but also reflects interorganelle communication mediated by functional protein interactions under conditions of stress.

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The authors declare no competing interest.

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Fig. 1. The Interaction between OsHAK21 and OsCYB5-2. (A) OsHAK21 interacts with OsCYB5-2 in yeast split-ubiquitin system. The 1:10 serial dilutions of yeast cells were spotted on control medium (Left) or selective medium (Middle). The same set of yeast transformants were assayed for β-gal activities developed on filter paper (Right). (B) Co-IP analysis for OsHAK21 and OsCYB5-2 interaction in agrobacterium-infiltrated tobacco leaves. OsHAK21-FLAG and HA-OsCYB5-2 were coexpressed as the indicated combinations (Top). **"-" represents vector alone. Proteins (Left) were immunoprecipitated with anti-HA antibody (IP: HA) and detected with anti-FLAG antibody (IB: FLAG). Proteins (Right) were immunoprecipitated with anti-FLAG antibody (IP: FLAG) and detected with anti HA antibody (IB: HA). Beads incubating proteins without (w/o) IgG are negative controls. (C) Rice protoplasts coexpressing OsHAK21-CFP and YFP-OsCYB5-2 were examined by confocal laser scanning microscopy. YFP-HDEL is a negative control of ER marker protein. (Scale bar, 20 μm.) (D) Quantitative FRET analysis for the interaction between OsHAK21-CFP and YFP-OsCYB5-2. The box plots depict the mean FRET efficiency from one experiment with n = 25 protoplasts. The boxes indicate the first and third quartiles, and the whiskers indicate the minimum to maximum values. The lines within the boxes indicate the median values. Statistically significant differences were determined by the two-tailed Student’s t test. Three independent experiments were repeated with similar results.

Results
High-Affinity K⁺ Transporter OsHAK21 Interacts with an ER-Localized OsCYB5. OsHAK21 has been identified as a functional K⁺/Na⁺ transporter that maintains K⁺/Na⁺ homeostasis in response to salt stress (8). To further elucidate the posttranslational regulatory mechanisms for OsHAK21, the yeast split-ubiquitin system was employed to screen for potential proteins that interact with OsHAK21 (31). Of 2.3 × 10⁶ transformants, we obtained 21 putative interactors, of which two CYB5-like heme/steroid binding domain–containing proteins (LOC_Os12g12170 [OsCYB5-1] and LOC_Os10g37420 [OsCYB5-2]) were identified (SI Appendix, Fig. S1). There are 14 members of the CYB5 family in rice and six members in Arabidopsis (SI Appendix, Fig. S2A) (26); we collectively named this family OsCYB5-n (n represents 1 through 14) (SI Appendix, Fig. S2A). OsCYB5-n were predicted to contain conserved structural characteristics: an N-terminal cytosolic heme-binding domain, a C-terminal transmembrane domain that anchors the protein to the ER or mitochondria/chloroplast, and a short luminal tail (SI Appendix, Fig. S2B) (21, 22). According to the β-glucuronidase (GUS) staining results, OsCYB5-1 showed weak transcription in young leaf. Roots of transgenic seedlings were not stained, while OsCYB5-2 expression was ubiquitous and strong in the tested tissues (SI Appendix, Fig. S3). We selected OsCYB5-2 as a putative OsHAK21 interactive partner for further analysis.

The interaction of OsHAK21–OsCYB5-2 was first determined using a yeast split-ubiquitin system (Fig. 1A). Yeast strains expressing the N-terminal half of yeast ubiquitin (NubWT) combined with the C-terminal half of yeast ubiquitin (Cub) fused with OsHAK21 (OsHAK21-Cub) were used as a positive control; yeast strains expressing Nub (G, mutant) combined with OsHAK21-Cub were used as a negative control. Yeast growth was observed for protein couples of positive control and OsHAK21-Cub/Nub-G-OsCYB5-2, suggesting that OsHAK21 and OsCYB5-2 physically interact. The results matched those obtained from a β-galactosidase (β-Gal) activity assay (Fig. 1A). To confirm the OsHAK21 and OsCYB5-2 interaction, C-terminally FLAG-tagged OsHAK21 and N-terminally hemagglutinin (HA)-tagged OsCYB5-2 were coexpressed in tobacco leaves, followed by immunoprecipitation (co-IP). The results showed that OsHAK21-FLAG and HA-OsCYB5-2 immunoprecipitated together (Fig. 1B).

To further investigate the interaction between OsHAK21 and OsCYB5-2 in plant cells, we first determined their cellular location. OsCYB5-2 was fused with green fluorescence protein (GFP) to produce GFP-OsCYB5-2; GFP-OsCYB5-2 showed a reticular morphology, which overlapped with the red fluorescence from mCherry-fused ER marker HDEL (SI Appendix, Fig. S4), indicating that OsCYB5-2 localized to the ER in plant cells. In previous work, OsHAK21 was shown to localize to the
As black triangle of yeast cells were spotted on the AP medium. (WT, athak5 transformed into R5421 as a positive control. mutant. The seedlings were grown for 12 d on the medium containing K

Fig. 2. OsCYB5-2 increases K⁺ transport activity of OsHAK21 in yeasts and plants. (A) OsCYB5-2 enhances the growth of K⁺-uptake-deficient yeast mutant R5421 expressing OsHAK21 on low-K⁺ AP medium. The WT yeast strain R757 was used as a positive control. OsAKT1, a K⁺ channel from Arabidopsis, was transformed into R5421 as a positive control. OsHAK21 and OsCYB5-2 were constructed into p416 and p424 vector, respectively. The 1:10 serial dilutions (as black triangle) of yeast cells were spotted on the AP medium. (B) Phenotypes of Arabidopsis transgenic lines expressing OsCYB5-2 and OsHAK21 in athak5 mutant. The seedlings were grown for 12 d on the medium containing K⁺ as the concentrations indicated. (C) Kinetics of Rb⁺ uptake in roots of WT, athak5, and Arabidopsis transgenic lines as described in B after 7 d of K⁺ starvation. The data are shown as means ± SD from n = 5 seedlings. Curves represent results of fitting Michaelis–Menten equation. FW, Fresh weight.

PM (8). We then applied a Förster resonance energy transfer (FRET) assay using OsCYB5-2 fused to yellow fluorescent protein (YFP-OsCYB5-2) and OsHAK21 fused to cyan fluorescent protein (OsHAK21-CFP). When YFP-OsCYB5-2 and OsHAK21-CFP were coexpressed in rice protoplasts, they produced high levels of FRET, which were 4.5-fold higher than the FRET generated during OsHAK21-CFP and YFP-HDEL coexpression (Fig. 1 C and D). Taken together, the results suggest specific binding between OsHAK21 and OsCYB5-2 in vivo.

In addition, the transgenic plants carrying OsCYB5-2 promoter::GUS showed that OsCYB5-2 was ubiquitously expressed in all tissues (SI Appendix, Fig. S3), and a similar pattern was found for OsHAK21 (8). A cross-section of GUS-stained roots showed strong signals in most cell types, consistent with the expression of OsHAK21 in xylem parenchyma and endodermal cells (SI Appendix, Fig. S3E) (8). Strong GUS activity driven by the OsCYB5-2 promoter was detected in germinating embryos (SI Appendix, Fig. S3I and J), similar to OsHAK21 expression during germination (17). These results suggest that the expression patterns of OsCYB5-2 and OsHAK21 are spatially and temporally similar, which increases the likelihood of interaction between OsHAK21 and OsCYB5-2 in rice.

OsCYB5-2 Enhances K⁺ Transport Activity of OsHAK21. To explore the biological significance of the interaction between OsCYB5-2 and OsHAK21, we examined how OsCYB5-2 expression affects OsHAK21 activity in the yeast heterologous expression system. When both OsCYB5-2 and OsHAK21 were coexpressed in yeast cells, the K⁺ uptake activity of OsHAK21 was improved by about 4.5-fold as compared with OsHAK21 expression alone at 0.5 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells. OsHAK21 activity was, however, weaker than that of the Arabidopsis K⁺ transporter AtKAT1 (34) and the WT yeast transporter R757. Coexpression of OsCYB5-2 and OsHAK21 further improved yeast growth under as low as 1 mM K⁺, suggesting that OsHAK21 exhibits K⁺-uptake activity in yeast cells.
OsHAK21 improved K⁺ uptake in Arabidopsis (Fig. 2B and SI Appendix, Fig. S5 B and C) (8, 35). The simultaneous expression of OsHAK21 and OsCYB5-2 in the athak5 mutant (athak5/OsHAK21/OsCYB5-2) improved plant growth even further, exhibiting increased root length and fresh weight compared to the athak5/OsHAK21 and WT plants. No significant changes in growth were observed when coexpressing OsCYB5-2 (Fig. 2B and SI Appendix, Fig. S5 B and C).

Direct measurements of K⁺-tracer Rb⁺ transport kinetics in plants revealed that overexpression of OsHAK21 complemented the impairment of HAK uptake in athak5 (Fig. 2C). Coexpression of OsCYB5-2 with OsHAK21 improved K⁺ uptake in plants compared to that of OsHAK21 only by increasing V_max and decreasing K_m. By contrast, overexpression of OsCYB5-2 only did not change the kinetic parameters for K⁺ uptake (SI Appendix, Fig. S5D). Together, these results indicate that OsCYB5-2 can increase OsHAK21 activity, thereby indirectly promoting K⁺ uptake in plants.

OsCYB5-2 and OsHAK21 Interaction Improves Salt Tolerance in Rice.
To test our hypothesis that interaction of OsCYB5-2 and OsHAK21 improves salt-stress tolerance, we first expressed OsCYB5-2 and OsHAK21 in a heterologous yeast system to examine its effect on growth at various NaCl concentrations. Yeast transformants expressing OsHAK21 or OsCYB5-2 could not grow vigorously at all NaCl concentrations (100 to 400 mM) tested. The combined expression of OsHAK21 and OsCYB5-2 significantly improved yeast growth, even at high (300 mM)-NaCl concentrations (SI Appendix, Fig. S6A). The improvement of salt tolerance by the combined overexpression of OsHAK21 and OsCYB5-2 was confirmed in transgenic Arabidopsis plants (SI Appendix, Fig. S6 B and C).

The interaction between OsHAK21 and OsCYB5-2 was then investigated in rice plants. OsCYB5-2 expression increased under salt stress, similar to that of OsHAK21 (SI Appendix, Fig. S7) (8). The OsCYB5-2-overexpressing rice plants with WT background (WT/OsCYB5-2-OE) showed high tolerance to salt stress and significantly higher fresh weight and chlorophyll content relative to WT plants transformed with empty vector (WT/vector) (Fig. 3 A–C). Moreover, when OsCYB5-2 was overexpressed in the oshak21 mutant background (8), no mitigating effects were observed (Fig. 3 A–C), suggesting that the function of OsCYB5-2 is OsHAK21 dependent.

To investigate whether the OsHAK21–OsCYB5-2 interaction regulates K⁺ and Na⁺ homeostasis in rice plants, their contents in the transgenic plants were analyzed. Under control conditions, no significant difference in Na⁺ (or K⁺) content or ratio was observed among the transgenic lines (Fig. 3 D–F and SI Appendix, Fig. S8). Following NaCl treatment for 12 d, WT/ OsCYB5-2-OE plants accumulated the lowest Na⁺ and highest K⁺ among the transgenic rice lines in both shoots and roots (Fig. 3 D and E and SI Appendix, Fig. S8 A and B), which resulted in the lowest Na⁺/K⁺ ratios (Fig. 3F and SI Appendix, Fig. S8C). Moreover, overexpression of OsCYB5-2 increased K⁺ net uptake and decreased Na⁺ net uptake under NaCl stress conditions (Fig. 3 G and H). Taken together, these results indicate that OsCYB5-2 increases OsHAK21 activity and promotes K⁺ uptake, which is essential for the maintenance of K⁺/Na⁺ homeostasis and salt tolerance in rice.

Salt Stress Triggers the OsHAK21–OsCYB5-2 Interaction.
We investigated whether and how salt stress affects the interaction between OsHAK21 and OsCYB5-2. We first used the yeast split-ubiquitin system to quantify the OsHAK21–OsCYB5-2 interaction (estimated based on the β-Gal activity; SI Appendix, Fig. S9A) and found that high Na⁺ significantly enhanced β-Gal activity in a dose- and time-dependent manner (SI Appendix, Fig. S9 B and N). We used OsHAK21-Cub+NubWT, which shows high β-Gal activity, as a control and found that the activity did not change at different concentrations of NaCl (0 to 400 mM) over 4 h. Another control, OsHAK21-Cub+NubG, also did not change according to the concentration of NaCl. The results suggest that the increase in β-Gal activity is specific for OsHAK21 and OsCYB5-2 binding. Importantly, the interaction did not vary according to the isotonic concentrations of K⁺ and mannitol or K⁺ deficiency (SI Appendix, Fig. S9). The results suggest that the increase in the degree of OsHAK21–OsCYB5-2 interaction is a specific response to high-Na⁺ stress.

To examine the OsHAK21–OsCYB5-2 interaction in rice cells, we developed constructs that enable coexpression of multiple chimeric fluorescent fusion proteins in suspensions cells (Fig. 4A and SI Appendix, Fig. S10 A and B) (36). The vectors facilitate fusion of the gene of interest with 3XFLAG-tagged CFP (FC) and HA-tagged YFP (YH), thus enabling detection of protein interactions using FRET and co-IP analysis (Fig. 4 A–D). We coexpressed OsHAK21-FC with YH-OsCYB5-2 in rice suspension cells of the oshak21 background. Transformant protoplasts were isolated to examine the OsHAK21–OsCYB5-2 interaction via FRET (Fig. 4 A and B). The resulting FRET efficiency, indicative of the OsHAK21–OsCYB5-2 interaction, was determined by dividing the emission intensity of FRET by the emission intensity of CFP (FRET/CFP) at predefined time points (37). The FRET efficiency (FRET/CFP) is proportional to the intensity of the two-protein interaction. Protein only expressing OsHAK21-FC and YH-OsCYB5-2 exhibited an increase in FRET efficiency following treatment with 100 mM NaCl but not with isotonic concentrations of mannitol (200 mM), indicating that the interaction between the two proteins was enhanced under salt stress (Fig. 4 B and C). NaCl treatment did not increase the interaction between another pair of proteins, AtVST1 in the peripheral PM and AtSR2C in the ER (SI Appendix, Fig. S10 A–D) (38); the interaction of these proteins has been shown to regulate stomatal development signaling (38). FRET efficiency changed in response to the addition of the bacterial flagellar peptide (flg22) to the protoplast expressing the flg22 receptor AtFLS2 and a receptor-like kinase (AtNIK1 or AtBIK1) (39, 40). However, the AtFLS2–AtNIK1/AtBIK1 interaction were not affected by NaCl or mannitol treatment (SI Appendix, Fig. S10 C–E). These results show that high-salt conditions specifically induce the interaction of OsHAK21 and OsCYB5-2 through ionic stress.

Suspension cells coexpressing OsHAK21-FC and YH-OsCYB5-2 were incubated in 100 mM NaCl, and the YH-OsCYB5-2/OsHAK21-FC interaction was quantified by performing co-IP over a time course of 60 min. The expression levels of OsHAK21-FC and YH-OsCYB5-2/OsHAK21-FC interaction did not change from 0 to 60 min of NaCl (0 or 100 mM) treatment. YH-OsCYB5-2/OsHAK21-FC binding increased following treatment with 100 mM NaCl, but binding did not change with 0 mM NaCl treatment (Fig. 4D and SI Appendix, Fig. S10F), suggesting that salt stress induces OsCYB5-2 binding to OsHAK21.

The K⁺ and Na⁺ contents were determined in rice suspension cells (oshak21 background) expressing either OsHAK21 (vector iii), OsCYB5-2 (vector iv), or both (vector ii) (Fig. 4A); expression was confirmed by transcription analysis (Fig. 4 F and G, Insets). Cells coexpressing OsCYB5-2 and OsHAK21 displayed increased K⁺ content and reduced Na⁺ accumulation at 90 to 120 min relative to transformants expressing OsHAK21 only incubated in salt (Fig. 4 E–G). The results suggest that salt stimulation triggers OsCYB5-2 binding to OsHAK21, which then mediates K⁺/Na⁺ homeostasis in cells; this is consistent with the genetic and physiological results (Fig. 3).

Leucine 128 in OsHAK21 Is a Key Residue for OsCYB5-2 Binding.
To identify the region of the OsHAK21 protein involved in OsCYB5-2 binding, serial deletion mutants of OsHAK21 were
constructed and tested in the yeast split-ubiquitin system (Fig. 5A). The cytoplasmic C-terminal fragment of OsHAK21 did not bind OsCYB5-2 (Fig. 5A). The C-terminal deletions up to 183-amino acid (aa) residues did not significantly affect OsCYB5-2 binding (Fig. 5A), suggesting that the OsCYB5-2 binding domain resides within the first 183-aa residues. To establish the essential residues for OsCYB5-2 binding within the first 183 residues, site mutations were made. In yeast systems, leucine (L) residues are thought to be essential for the binding of sugar (and sorbitol) transport proteins with MdCYB5 from apple plants (29). We therefore performed site-directed mutagenesis to separately replace each of the 10 L residues (within OsHAK21) with alanine (A). The Na+ content (mmol g DW-1) of Shoots under NaCl treatment was measured (Fig. 5B). A Student’s t-test was used to determine the significance of the differences.

Fig. 3. OsCYB5-2 improves salt tolerance in rice by regulating OsHAK21-mediated K⁺ transport. (A–C) Phenotypes of OsCYB5-2-overexpressed lines in WT (Nipponbare) and oshak21 backgrounds. Rice seedlings were hydroponically grown with or without 150 mM NaCl for 12 d. Representative photographs of plants (A), total chlorophyll in shoots (B), and fresh weight (C) are shown. The transformed empty vector (pCM1307) seedlings were used as negative controls. (D–F) Effects of OsCYB5-2-overexpression on Na⁺ and K⁺ accumulation in shoots under salt stress. Seedlings were treated as in A, and the shoots were harvested for Na⁺ and K⁺ content assay. Data are shown as means ± SD (B and C, n = 12; D–F, n = 5 biologically independent seedlings for each transgenic rice lines). Lowercase letters above the bars in B–F indicate significant differences among means (P value = 0.05, Kruskal–Wallis bilateral test). ns indicates nonsubstantial differences at that level of significance. (G and H) K⁺ and Na⁺ net uptake rates in rice seedlings during 10 d of the treatment with 150 mM NaCl. Data in G and H are shown as means ± SD (n = 5). Statistically significant differences were determined by the two-tailed Student’s t test.
Fig. 4. The interaction between OsHAK21 and OsCYB5-2 is triggered by salt treatment. (A) Schematic diagram of the coexpression proteins integrated into a vector. The vectors (i and ii) feature two independent expression cassettes with two strong promoters (2×35S and 3S promoter) for high-level expression in plant cells. The vector (i) encoding CFP and YFP fusion proteins also encoded an in-frame 3×FLAG and HA tag as indicated and consequently were dual-use for FRET (iii and iv). Proteins coexpressed from ii without fusion tag were used to detect the function of ion transport (E–G). iii and iv were used as controls for protein expression alone. NosT, terminator of the Nos gene. (B) The FRET efficiency (FRET/CFP) of the interaction triggered by 100 mM NaCl and 200 mM mannitol in protoplasts coexpressing OsHAK21-FC+YH-OsCYB5-2 (i in A), FC, CFP-CFP Tag; YH, YFP-HA Tag. The arrow indicates the addition of treatments. The data represent means ± SD from the determination of n = 10 rice protoplasts for each treatment. Three independent experiments were repeated with similar results. (C) Representative FRET images of cells from B. (Scale bar, 20 μm.) (D) Time-lapse co-IP assay of the interaction between OsHAK21-FC and YH-OsCYB5-2 (i in A) in oshak21 suspension cells treated with 100 mM NaCl. The same quality of proteins (5 μg) from different time points were immunoprecipitated with anti-FLAG beads (IP: FLAG) and detected with anti-HA antibody (IB: HA). The experiment was performed independently three times, and representative results are shown. Bands relative values were determined by ImageJ software. The relative protein level at each time point was normalized to OsHAK21-FC of input, and the value at 0 min was set as standard 1. (E–G) Time-course accumulation of K⁺ content (E), Na⁺ content (F), and Na⁺/K⁺ ratio (G) in oshak21 suspension cells expressing protein combinations (ii through iv in A) with 100 mM NaCl treatments in the presence of 1 mM KCl. Insets show the transcripts of OsHAK21 (F) and OsCYB5-2 (G) in independent oshak21 suspension cells expressing protein combinations as indicated with different colors in E. The data are shown as means ± SD from n = 5 biologically suspension cells lines for each protein combination. Statistically significant differences were determined by the two-tailed Student’s t test. Three independent experiments were done with similar results.
the identified 1- to 183-aa fragment) with proline (P) in full-length OsHAK21. Finally, the L128P mutation, which lies in the intracellular loop region between transmembrane regions 2 and 3, disrupted OsCYB5-2 binding (Fig. S4 and SI Appendix, Fig. S11 A and B). The L128P mutation did not change the expression or PM localization of OsHAK21 (SI Appendix, Fig. S11 C–G). Mutation of other L residues did not significantly influence OsCYB5-2/OsHAK21 binding (SI Appendix, Fig. S11 A and B). The assay was repeated in tobacco leaves using luciferase complementation imaging (LCI) and co-IP (Fig. 5B and SI Appendix, Fig. S11E), which confirmed the yeast split-ubiquitin results. It is worth noting that L128 of OsHAK21 is conserved among representative HAK family members in different plant species (Fig. 5C).

To further reveal the role of OsCYB5-2 binding in K⁺ transport mediated by OsHAK21, a kinetic characterization of Rb⁺ (K⁺) transport was performed in yeast cells. Coexpression of OsCYB5-2 together with OsHAK21 increased the affinity for

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**Fig. 5.** OsCYB5-2 interacts with OsHAK21 at L128. (A) The interaction of different OsHAK21 truncations and OsCYB5-2. In the schematic structures of OsHAK21 (left), the gray portions indicate the sequences removed. The asterisk indicates OsHAK21 with the point mutation at L128P. The right panels reveal the interaction between different portions of OsHAK21-nLuc and cLuc-OsCYB5-2 in tobacco leaves. The cLuc-RAR1 served in HAK families of different species. (B) Initial rates of Rb⁺ uptake in yeast R5421 strain expressing different gene combinations. Curves represent results of fitting Michaelis–Menten equation. Kₘ and Vₘₐₓ values are shown in SI Appendix, Fig. S11F. DW, dry weight. Three independent experiments were carried out, and the data represent the mean ± SD from n = 3 biologically independent yeast lines for each genotype. (E and F) Growth curves of the R5421 strain transformed with OsHAK21 and mutational OsHAK21 with OsCYB5-2 in liquid AP medium supplemented with 10 mM K⁺ (E) and 0.5 mM K⁺ (F). Note that the curves of OsHAK21, OsHAK21L128P, and OsHAK21L128P+OsCYB5-2 almost overlap in D–F.
OsHAK21 directly (Fig. 6A). We measured the apparent dissociation constant \( K_d \) of K\(^+\) and OsHAK21 using isothermal titration calorimetry (ITC). As direct binding measurements of transporters and substrates can be difficult because of low substrate affinity and low levels of purified protein (41), we expressed full-length OsHAK21 protein in Spodoptera frugiperda 9 insect cells and purified the protein (SI Appendix, Fig. S12A). ITC was performed by titrating a solution containing K\(^+\) into an ITC chamber, with OsHAK21 protein dissolved in buffer with 50 mM NaCl as the background electrolyte for solubilization (Fig. 6A, Top). The heat from each injection was used to obtain the apparent \( K_d \) of 1.36 mM (Fig. 6A, Bottom). When 50 mM lithium chloride (LiCl) was used as the background electrolyte, similar \( K_d \) values were recorded.

OsCYB5-2 Increases Apparent Affinity of OsHAK21 for K\(^+\)-binding. To investigate the biochemical mechanisms by which OsCYB5-2 improves OsHAK21-mediated K\(^+\) transport, we measured the apparent dissociation constant \( K_d \) of K\(^+\) and OsHAK21 using isothermal titration calorimetry (ITC). As direct binding measurements of transporters and substrates can be difficult because of low substrate affinity and low levels of purified protein (41), we expressed full-length OsHAK21 protein in Spodoptera frugiperda 9 insect cells and purified the protein (SI Appendix, Fig. S12A). ITC was performed by titrating a solution containing K\(^+\) into an ITC chamber, with OsHAK21 protein dissolved in buffer with 50 mM NaCl as the background electrolyte for solubilization (Fig. 6A, Top). The heat from each injection was used to obtain the apparent \( K_d \) of 1.36 mM (Fig. 6A, Bottom). When 50 mM lithium chloride (LiCl) was used as the background electrolyte, similar \( K_d \) values were recorded (SI Appendix, Fig. S13A).

Fig. 6. OsCYB5-2 increases the apparent affinity of OsHAK21 for K\(^+\)-binding. (A–C) ITC profiles and thermodynamic data of OsHAK21 (A), OsHAK21+OsCYB5-2ΔC (B), and OsHAK21+apo-OsCYB5-2ΔC (C) titrated with K\(^+\). In total, 19 injections of KCl solution were added to protein solution in ITC chamber. During each injection, a small amount of KCl is rapidly mixed with the protein, from which heat is exchanged and recorded in the resulting thermogram. The area of each injection peak (Top of A–C) is equal to the heat released from that injection with time. The complete binding isotherm for K\(^+\)-protein interaction (Bottom of A–C) was obtained by integrated heat plotted against the molar ratio (ligand/protein) injection of K\(^+\) and OsHAK21 using SI Appendix, Fig. S13). OsHAK21+OsCYB5-2ΔC and OsHAK21+apo-OsCYB5-2ΔC. (E) Biolayer interferometry (BLI) analysis for the interactions between OsHAK21 and OsCYB5-2ΔC and OsHAK21 and apo-OsCYB5-2ΔC.

OsCYB5-2 reduces OsHAK21 sensitivity to Na\(^+\). In saline environments, plant cells accumulate high concentrations of Na\(^+\), which prompted us to investigate whether high-salt concentrations affect OsHAK21 affinity for K\(^+\). We measured the apparent K\(^+\) affinity in the presence of different concentrations of NaCl. As seen in Fig. 7A, NaCl concentrations (50 to 200 mM) reduced the affinity of OsHAK21 for K\(^+\) by increasing the \( K_d \), and the reduction was dose dependent. As Na\(^+\) does not bind OsHAK21 directly (SI Appendix, Fig. S13B), the reduction in apparent affinity for K\(^+\) could have been caused by the high

Affinity in the presence of different concentrations of NaCl. As seen in Fig. 7A, NaCl concentrations (50 to 200 mM) reduced the affinity of OsHAK21 for K\(^+\) by increasing the \( K_d \), and the reduction was dose dependent. As Na\(^+\) does not bind OsHAK21 directly (SI Appendix, Fig. S13B), the reduction in apparent affinity for K\(^+\) could have been caused by the high
When OsCYB5-2ΔC was added to the solution, the reduction in OsHAK21 apparent affinity for K\(^{+}\) was significantly less pronounced at all NaCl concentrations examined (Fig. 7A); this effect was not observed with added apo-OsCYB5-2ΔC. Furthermore, NaCl increased the binding affinities between OsHAK21 and OsCYB5-2ΔC, as determined by ITC determination (Fig. 7B). BLI analysis for the interaction between OsHAK21 and OsCYB5-2ΔC at different Na\(^{+}\) concentrations in solution (Fig. 7C–E) showed that NaCl enhanced the association rate and dissociation rate of OsHAK21 and OsCYB5-2ΔC. Lineweaver–Burk double-reciprocal plots for Rb\(^{+}\) uptake in yeast expressing OsHAK21 (Figs. 7C and 7D) and OsHAK21+OsCYB5-2 (Figs. 7D and 7E) in the absence (0 mM) or presence of 10 or 50 mM Na\(^{+}\). Na\(^{+}\) K\(_{i}\) represents the inhibition constant of Na\(^{+}\). DW, dry weight. All experiments have been repeated three times, and the data are shown as mean ± SD (n = 3). (F) Schematic model for OsCYB5-2 and OsHAK21 interaction in salt response. Salt stress enhances ER-localized OsCYB5-2 binding to PM-localized OsHAK21, promoting OsHAK21 affinity and preference for K\(^{+}\)-binding. As a result, OsHAK21-mediated, inward K\(^{+}\) transport maintains intracellular K\(^{+}\)/Na\(^{+}\) homeostasis and ultimately improves salt tolerance in rice.

Ionic strength of the solution. When OsCYB5-2ΔC was added to the solution, the reduction in OsHAK21 apparent affinity for K\(^{+}\) was significantly less pronounced at all NaCl concentrations.
using BLI methods with biotin-labeled proteins (Fig. 7B), consistent with the FRET results (Fig. 4B). Importantly, OsHAK21 and OsCYB5-2ΔC bind at a physiologically viable level (nanoliter), suggesting that the binding could occur in plant cells.

To functionally characterize the affinity of OsCYB5-2-OsHAK21 for K+ under salt treatment, kinetic parameters (inhibition constant $K_i$ for Na+) were assessed in yeast cells. The $R_b(K^+)$-uptake in the presence of Na+ demonstrated that Na+ resulted in competitive inhibition, with a $K_i$ of 18.17 mM for $R_b(K^+)$-uptake in cells expressing OsHAK21 (Fig. 7C). The $K_i$ of Na+ was increased 2.6-fold by the expression of OsCYB5-2 and OsHAK21 compared to OsHAK21 alone (Fig. 7C and D), suggesting that OsCYB5-2 alleviated the inhibitory effect of Na+ on OsHAK21. The L128P mutation did not obviously change the inhibition of OsHAK21 by Na+ but abolished the allelic effects of OsCYB5-2 on OsHAK21 (Fig. 7C and SI Appendix, Figs. S11 F and J).

To explore the effect of the electron carrier properties of OsCYB5-2 on OsHAK21-mediated K+-uptake, we generated OsCYB5-2mut by substituting two conserved His residues with alanine (H40A/H64A) to impair the coordination with heme iron and the electron transfer properties of OsCYB5-2 (SI Appendix, Fig. S14A) (24, 26). Like the L128P mutation in OsHAK21, OsCYB5-2mut was unable to stimulate the transport activity of OsHAK21 (SI Appendix, Figs. S11H and 14B) and recovered the inhibitory effect of Na+ on OsHAK21-mediated $R_b(K^+)$-uptake (Fig. 7C and E). However, mutation of OsCYB5-2mut did not change its association with OsHAK21 or ER localization (SI Appendix, Fig. S14 C–G). Taken together, these findings demonstrate that heme-binding and thus the electron transfer properties of OsCYB5-2 are essential for regulating the transport activity of OsHAK21 by improving K+-binding, especially under NaCl stress.

**Discussion**

Our understanding of effective quantitative trait loci, genes, and pathways that play roles in the avoidance of Na+ toxicity at cellular and tissue levels has steadily improved (45–47). Evidence is also increasing regarding the importance of K+-uptake (via HAKs, AKTs, and HKTs, etc.) and K+/Na+ homeostasis under salt stress (4, 47, 48), although no mechanistic insights into salt-related regulation of K+ transporter have been achieved. In this study, we report a posttranslational mechanism for the regulation of HAK transporter activity by ER-localized OsCYB5-2. This salt-triggered mechanism contrasts the interference of Na+ with K+ high-affinity transport and therefore plays an important role in maintaining K+/Na+ homeostasis under salt stress in plants.

Cellular adaptation to stressful environments requires coordinated, interorganellar responses to transduce stress signals and maintain the integrity of cellular structures in both animal and plant cells (49). In a previous study, a functional connection between the PM and microtubules (MTs) was discovered, whereby lipid phosphatidic acid binds to MT-associated protein 65 in response to salt stress (50). More recently, lipid-associated SYT1 contact site expansion in Arabidopsis under salt stress was reported, resulting in enhanced ER–PM connectivity (49). However, the role of ER–PM connection in stress adaptation remains unclear. Here, we report that salt stress triggers a rapid ER–PM connection via binding of ER-localized OsCYB5-2 and PM-localized OsHAK21. OsCYB5-2 and OsHAK21 binding and hence ER–PM connection occurred as quickly as 50 s after the onset of NaCl treatment (Fig. 4), which is quicker than that in Arabidopsis, in which phosphoinositide-associated SYT1 contact site expansion occurs within hours (49).

OsCYB5-2 and OsHAK21 interaction was not only observed at the protoplast and cellular level (Figs. 1 and 4) but also in whole rice plants. Overexpression of OsCYB5-2 conferred increased salt tolerance to WT plants but not to oshak21 mutant plants that lack the partner protein OsHAK21 (Fig. 3), providing further evidence that the OsCYB5-2–OsHAK21 interaction plays a positive role in regulating salt tolerance.

Plant HAK transporters are predicted to contain 10 to 14 transmembrane domains, with both the N and C termini facing the cytoplasm (51). On the N-terminal side, the GD(EGTF)-FALY motif is highly conserved among members of the HAK family (Fig. 5C) (52). The L128 residue, which is required for OsCYB5-2 binding, is located within the GDGGGTFAFY motif (Fig. 5). Residue substitution (F130S) in AtHAK5 led to an increase in K+ affinity by >100-fold in yeast (52). AtHAK5 activity was also found to be regulated by CIPK23/CBL1 complex–mediated phosphorylation of the N-terminal 1- to 95-aa residues (14). In rice, a receptor-like kinase RUPO interacts with the C-tail of OsHAKs to mediate K+ homeostasis (53). Thus, the L128 bound by OsCYB5 identified in this work is uniquely involved in HAK transporter regulation.

OsCYB5-2 binding at L128 elicits an increase in K+-uptake (Fig. 5D), consistent with the role of OsCYB5-2 in enhancing the apparent affinity of OsHAK21 for K+ (binding) (Fig. 6). An important question is raised by this: how does OsCYB5-2 regulate OsHAK21 affinity for K+? Electron transfer between CYB5 and its redox partners is reliant upon its heme cofactor (24, 42). Given that both apo-OsCYB5-2ΔC (no heme) and OsCYB5-2mut are unable to stimulate K+ affinity of OsHAK21 (Figs. 6 and 7 and SI Appendix, Figs. S14 and S15), we propose that electron transfer is an essential mechanism for OsCYB5-2 function. This could occur via redox modification of OsHAK21 to increase K+ affinity. We cannot, however, rule out the possibility of allosteric effects of OsCYB5-2 binding on OsHAK21. Several residues in AtHAK5 have been proposed as the sites of K+-binding or -filtering (20, 54). Following association of OsCYB5-2 with residue L128 of OsHAK21, a conformational change likely occurs in OsHAK21, resulting in a modulated binding efficiency for K+.

Active transporters and ion channels coordinate to produce and dissipate ionic gradients, allowing cells to control and finely tune their internal ionic composition (55). However, under salt stress, apoplastic Na+ entry into cells depolarizes the PM, making channel-mediated K+-uptake thermodynamically impossible. By contrast, activation of the gated, outward-rectifying K+ channel induces K+ efflux out of cells. Together, these effects dramatically reduce the K+ concentration in plant cells. K+-uptake is therefore dependent on active transport via K+/H+ symport mechanisms (HAK family), which are driven by the proton motive force generated by H+-ATPase (48). A strong, positive correlation between H+-ATPase activity and salinity stress tolerance has been reported (56, 57). In rice, OsHAK21 is essential for salt tolerance at the seedling and germination stages (8, 17). OsHAK21-mediated K+-uptake increased with lowering of the external pH (increasing H+ concentration); this effect was abolished in the presence of the proton ionophore FCCP (SI Appendix, Fig. S15 D–F). Confirmation of synergistic effects of oxidoreduction and H+ concentration on OsHAK21 activity requires further study. The CYB5-mediated OsHAK21 activation mechanism reported here differs from the posttranslational modifications that occur via phosphorylation by the CBL/CIPK pair (11, 19, 20), which likely relies on salt perception (which triggers calcium signals) (58).

We propose that salt triggers association of ER-localized OsCYB5-2 with PM-localized OsHAK21, causing the OsHAK21 transporter to specifically and effectively capture K+. As a result,
OsHAK21 transports K⁺ inward to maintain intracellular K⁺/Na⁺ homeostasis, thus improving salt tolerance in rice (Fig. 7F).

Materials and Methods

Information on plant materials used, growth conditions, and experimental methods employed in this study is detailed in **SI Appendix**. The methods include the methods on vector construction and plant transformation, co-IP assay, FRET analysis, subcellular localization, yeast two-hybrid, histochemical staining, gene expression analysis, LCI assay, BBI plant treatment, and ion content determination. Details of experimental conditions for ITC are provided in **SI Appendix, Table S1**. Primers used in this study are listed in **SI Appendix, Table S2**.

Data Availability. All study data are included in the article and/or **SI Appendix**.

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