

Structural insights into the negative regulation of BRI1 signaling by BRI1-interacting protein BKI1

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Brassinosteroids (BRs) are essential steroid hormones that have crucial roles in plant growth and development. BRs are perceived by the cell-surface receptor-like kinase brassinosteroid insensitive 1 (BRI1). In the absence of BRs, the cytosolic kinase domain (KD) of BRI1 is inhibited by its auto-inhibitory carboxyl terminus, as well as by interacting with an inhibitor protein, BRI1 kinase inhibitor 1 (BK11). How BR binding to the extracellular domain of BRI1 leads to activation of the KD and dissociation of BK11 into the cytosol remains unclear. Here we report the crystal structure of BRI1 KD in complex with the interacting peptide derived from BK11. We also provide biochemical evidence that BRI1-associated kinase 1 (BAK1) plays an essential role in initiating BR signaling. Steroid-dependent heterodimerization of BRI1 and BAK1 ectodomains brings their cytoplasmic KDs in the right orientation for competing with BK11 and transphosphorylation.

Keywords: BR signaling; BK11 inhibition; BRI1 activation; BRI1-BAK1 transphosphorylation

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Introduction

Signal perception through cell surface receptors, and transduction of this signal to the cell's interior, are essential to all life forms [1]. In plants, a large family of such receptors is the leucine-rich repeat receptor-like kinases (LRR-RLKs), with more than 220 members in *Arabidopsis* and about 400 in rice. Each LRR-RLK contains a leucine-rich repeat extracellular domain that usually binds ligands, a single membrane-spanning helix, and a cytoplasmic kinase domain that initiates intracellular signal

transduction [2]. Two of the well-studied LRR-RLKs are brassinosteroid-insensitive 1 (BRI1), the brassinosteroid (BR) receptor, and BRI1-associated kinase 1 (BAK1) that serves as a coreceptor for BRI1 activation [3-5]. BRs are plant-specific steroids that play essential roles in a wide range of developmental and physiological processes, such as stem elongation, leaf development, vascular differentiation, male fertility, flowering time, senescence, stomata development, and resistance to biotic and abiotic stresses [6-9]. Mutations of BRI1 or other components of the BR biosynthetic or signaling pathway cause severe growth abnormalities, including dwarfism, male sterility, delayed flowering, reduced apical dominance and photomorphogenesis in darkness. In contrast, increased BR content or enhanced BR signaling can enhance cellulose biosynthesis, biomass accumulation, and plant growth.

BR signaling is initiated by BR binding to a cell surface receptor kinase, BRI1 [10-12]. In the absence of BRs, the BRI1 KD is kept in a basal state by its auto-inhibitory carboxyl terminus [13] and by interacting with an inhibitor protein BK11 [14]. BR binding to the extracellular domain of BRI1 causes autophosphorylation of the BRI1 KD, release of BRI1 C-terminal autoinhibition,

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Abbreviations: BRs (brassinosteroids); BRI1 (brassinosteroid insensitive 1); BK11 (BRI1 kinase inhibitor 1); BAK1 (BRI1-associated kinase 1); LRR-RLKs (leucine-rich repeat receptor-like kinases); BIM (BRI1-interacting motif); CDs (cytosolic domains); KD (kinase domains); Cter (C-terminal); BSKs (BR-signaling kinases); JM (juxtamembrane); rmsd (root-mean-square deviation); PKA (protein kinase A); MAKR (membrane associated kinase regulators); PM (plasma membrane); pBK11-Cter (phosphorylated BK11-Cter)

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and BKI1 phosphorylation and its dissociation from the plasma membrane, leading to the formation of BRI1 and BAK1 complex [13, 15-17]. The KDs of BRI1 and BAK1 transphosphorylate each other on multiple sites to form fully activated receptor [17]. The activated BRI1 can phosphorylate several receptor-like cytoplasmic kinases, including the BR-signaling kinases (BSKs) and constitutive differential growth 1, which interact with and presumably activate the *bri1*-suppressor 1 phosphatase to transduce BR signaling downstream of BRI1 [18, 19]. The BSK family includes 12 members (BSK1-12) which play redundant roles in BR signaling [20].

Although the activation of BR signaling has been extensively studied during the past decade, a major gap in our understanding lies in the early molecular events that regulate BR-induced BRI1/BAK1 association and reciprocal phosphorylation. How BR binding to BRI1 ectodomain leads to partial activation or autophosphorylation of BRI1 KD has remained unclear. In the present study, we report the crystal structure of BRI1 KD in complex with the BRI1-interacting motif derived from BKI1. We also provide clear biochemical evidence that BAK1 plays an essential role in the early events of BRI1 activation, and BKI1 competitively inhibits transphosphorylation on the cytosolic domains of BAK1 and BRI1. Steroid-induced heterodimerization of BRI1 and BAK1 ectodomains brings their cytoplasmic KDs in the right orientation for competing with BKI1 and transphosphorylation. These results not only reveal the structural basis underlying BKI1 recognition by BRI1, but also provide insight into the initiating mechanism of brassinosteroid-induced BRI1 activation.

Results

Biochemical characteristics of the cytosolic domains of BRI1 and BAK1

To understand how BKI1 negatively regulates BRI1 signaling, we expressed and purified the cytosolic domains of BRI1 (BRI1-CD, residues 815-1196) and BAK1 (BAK1-CD, residues 255-615), the KD of BSK3 (BSK3-KD, residues 36-329), and the C-terminal region of BKI1 (BKI1-Cter, residues 257-337; Figure 1A). Bacterially expressed BRI1 and BAK1 proteins are catalytically active, because they undergo autophosphorylations during the expression and purification procedure [17, 21]. Western blotting confirmed that the recombinant BRI1-CD and BAK1-CD are phosphorylated on multiple sites when purified from *E. coli* (Figure 1B). The phosphorylated proteins can be dephosphorylated by a widely used protein phosphatase PP2C α , and the corresponding kinase-dead mutants, BRI1-CD^{K911R} (mBRI1-CD) and

BAK1-CD^{D434N} (mBAK1-CD), are completely unphosphorylated (Figure 1B). The purified mBAK1-KD, but not mBRI1-KD, can be rapidly phosphorylated by BRI1-CD in the presence of 1 mM ATP and 10 mM Mg²⁺ [21]. In contrast, mBRI1-CD, rather than mBAK1-CD, can be phosphorylated by BAK1-CD (Figure 1C). The inability of an active BRI1-CD to transphosphorylate an inactive mutant mBRI1-CD suggests that the mechanism of autophosphorylation is intramolecular or mediated by the nascent kinase passing through a transitory intermediate form [22].

To characterize the steady-state kinetic properties of BRI1-CD, the enzyme-catalyzed phosphorylation reaction was monitored using an enzyme-coupled spectrophotometric assay that couples production of ADP to oxidation of NADH, measured as a decrease in absorbance at 340 nm [23]. Figure 1D shows a typical progress curve of the BRI1-catalyzed BSK3-KD phosphorylation. When all of the substrates have been converted to products, the phosphorylation stoichiometry of BSK3-KD was determined to be close to 1 mol of phosphate/mol of BSK3-KD (Supplementary information, Figure S1), consistent with the previous findings that BRI1 specifically phosphorylated BSK3-KD at Ser212 within the BRI1 consensus phosphorylation sequence (²⁰⁶RDGKSYSTNLAFT²¹⁸) both *in vitro* and *in vivo* [18]. The BRI1-CD-catalyzed reaction obeyed classical Michaelis-Menten kinetics (inset of Figure 1D). Direct curve fitting of the data to the Michaelis-Menten equation yielded k_{cat} and K_m values, $3.99 \pm 0.21 \text{ s}^{-1}$ and $20.97 \pm 3.30 \mu\text{M}$, respectively, indicating that BSK3 is an efficient substrate for BRI1.

BKI1 negatively regulates BR signaling through two evolutionarily conserved motifs: a lysine/arginine-rich motif that targets BKI1 to the plasma membrane, and a BRI1-interacting motif (BIM, residues 306-325) that binds to the BRI1 KD. There are two phosphorylation sites, Ser270 and Ser274, in BKI1-Cter, which play important roles in subsequent phosphorylation of Tyr211 and dissociation of BKI1 from the plasma membrane [14, 15, 19]. *In vitro* kinase assay confirmed that the purified BKI1-Cter can be rapidly phosphorylated by BRI1-CD (Figure 1E). Inset of this figure shows the variation of initial rates of the BRI1-catalyzed reaction with BKI1-Cter concentrations. By fitting the Michaelis-Menten equation to the experimental data, the values of k_{cat} and K_m were determined to be $0.97 \pm 0.06 \text{ s}^{-1}$ and $4.28 \pm 0.73 \mu\text{M}$, respectively. Both the k_{cat} and K_m values decreased about fourfold as compared with those of BSK3-KD substrate, suggesting that the two substrates may have distinct interaction modes with BRI1-CD. The substrate specificity constant k_{cat}/K_m for BRI1-catalyzed BKI1-Cter phosphorylation was $2.3 \times 10^5 \text{ M}^{-1}\text{s}^{-1}$, which is similar

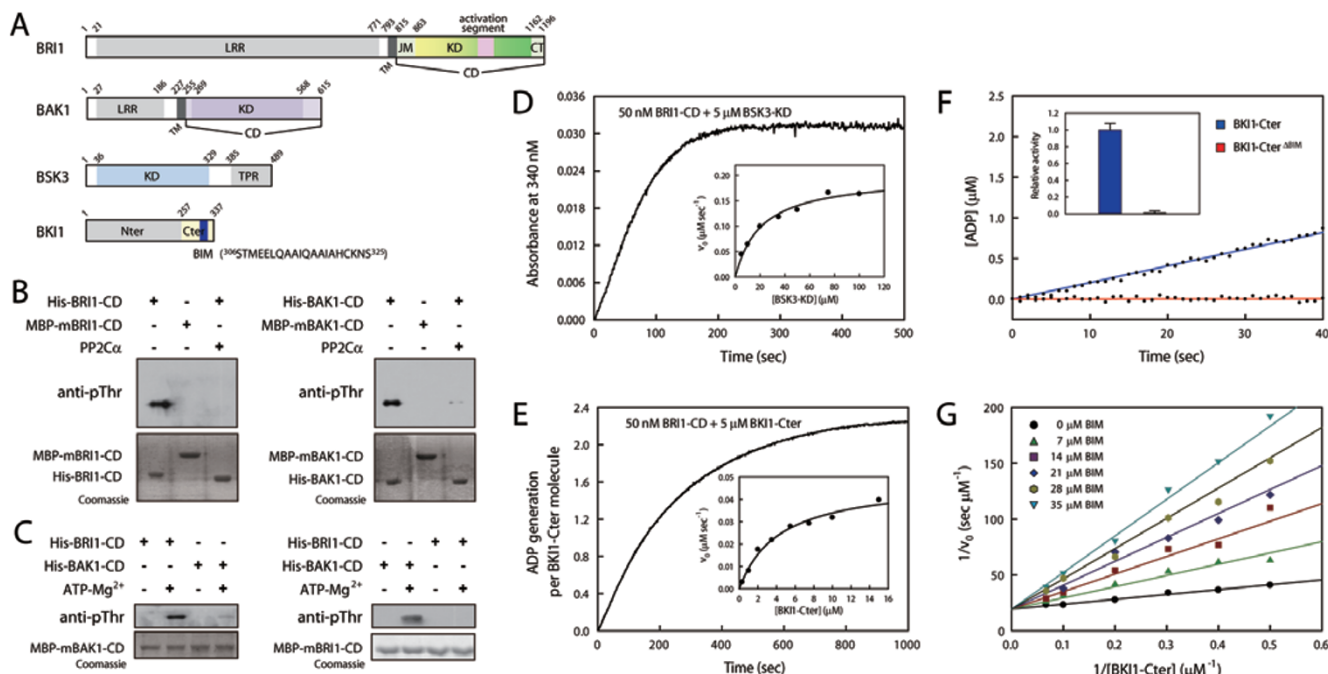


Figure 1 BIM is a major determinant for BRI1-catalyzed BKI1 phosphorylation. **(A)** Schematic diagram of BRI1, BAK1, BSK3 and BKI1. LRR, leucine-rich repeat domain; TM, transmembrane domain; CD, cytosolic domain; KD, kinase domain; TPR, tetratricopeptide repeat domain; Nter, N-terminal domain; Cter, C-terminal domain; BIM, BRI1-interacting motif. The sequence of the BIM peptide (blue) is provided. **(B)** Phosphorylation states of recombinant wild-type, kinase-dead mutant and PP2C α -dephosphorylated BRI1-CD and BAK1-CD analyzed using a general anti-pThr antibody. **(C)** Transphosphorylation analyses of BRI1-CD and BAK1-CD by western blotting. **(D and E)** Representative time course of BRI1-CD-catalyzed phosphorylation on BSK3-KD **(D)** and BKI1-Cter **(E)**. The insets show the initial rates vs the BSK3-KD and BKI1-Cter concentrations, respectively. **(F)** Time courses of BRI1-CD-catalyzed phosphorylation of BKI1-Cter and BKI1-Cter^{ΔBIM} (5 μ M). The relative activities of BRI1-CD towards two substrates (mean \pm SEM, $n = 3$) are shown in the inset. **(G)** Competitive inhibition of BRI1-catalyzed BKI1-Cter phosphorylation by BKI1-BIM. The kinetic parameters were obtained by fitting the experimental data to the competitive inhibition equation. See also Supplementary information, Data S1 and Figure S1.

to that of the BRI1-catalyzed BSK3-KD phosphorylation ($1.9 \times 10^5 \text{ M}^{-1}\text{s}^{-1}$). In comparison, neither BSK3-KD nor BKI1-Cter could be effectively phosphorylated by BAK1-CD under the same experimental conditions.

To quantitatively assess the contribution of the BIM peptide to the BRI1-catalyzed reaction, we measured the phosphorylation efficiency of BKI1-Cter^{ΔBIM} (a BIM-deleted BKI1-Cter mutant) by BRI1-CD. As shown in Figure 1F, compared with intact BKI1-Cter, deletion of the BIM region resulted in a dramatic drop (more than 80-fold) of the phosphorylation rate, suggesting that the phosphoacceptor sequence of BKI1-Cter in itself binds weakly to the active site of BRI1-CD, and the BRI1-interacting motif is a major determinant for the substrate specificity of BKI1-Cter [15, 19]. We also investigated the mechanism by which a synthetic BIM peptide (³⁰⁶STMEELQAAIQAAIAHCKNS³²⁵) inhibits BRI1 kinase activity towards BKI1-Cter substrate. The kinetics of the BRI1-CD-catalyzed phosphorylation of BKI1-Cter

were examined in the presence of various fixed concentrations of BIM peptide. Double-reciprocal plots of initial rate data over a range of concentrations for both the BIM peptide and BKI1-Cter substrate revealed a simple competitive pattern with a K_i value of $5.40 \pm 0.90 \mu\text{M}$ (Figure 1G), which is comparable to the K_m value for the BRI1-CD-catalyzed BKI1-Cter phosphorylation. This result further confirms that the BIM peptide was sufficient to tightly interact with the BRI1 KD [15].

Crystal structure of BRI1 KD in complex with BIM peptide

First, we carried out kinase activity assays with three fragments of the BRI1 cytoplasmic domain to assess the roles of the juxtamembrane (JM) and Cter regions (Supplementary information, Table S1). The JM region, KD and Cter segment have been previously assigned using a BRI1 homology model [15]. In the presence of the JM region, the K_m values of the fragments containing the JM

and/or Cter regions are only slightly lower than that of the KD alone. Our results are in contrast to those of Oh *et al.* [24] who reported that deletion of JM residues up to 882 caused a severe reduction in BRI1 kinase activity. One possible explanation for this discrepancy between the two studies might be that the construct of BRI1 KD used in their study was not fully active since deletion of residues 863-882 (the α B helix in the BRI1-KD structure) has a strong impact on solubility of the construct.

To gain a structural understanding of the mechanism

of BKII inhibition, we systematically screened for crystallizable fragments of the BRI1 cytoplasmic domain and obtained crystals of BRI1-KD (863-1162) in complex with the BKII-BIM peptide and AMP-PNP. The crystal structure of the ternary complex was determined at 2.2 Å resolution (Table 1). It should be indicated that the JM region is crucial for *in vivo* BRI1 function [13, 24, 25], and thus some aspects of BRI1's physiological function cannot be fully elucidated using our structural model. The crystallographic asymmetric unit contains two mol-

Table 1 The structural and refinement statistics for the BRI1-KD-BIM complex

	BRI1-KD-BIM (Native)	BRI1-KD-BIM (SeMet)
Data Collection^a		
Wavelength (Å)	0.9793	0.9789
Space group	$P2_1$	$P2_1$
Unit cell		
<i>a, b, c</i> (Å)	62.92, 80.14, 79.62	63.715, 80.288, 80.557
α, β, γ (°)	90.00, 108.86, 90.00	90.00, 109.55, 90.00
Resolution (Å)	50-2.25 (2.33-2.25) ^b	50-2.77 (2.87-2.77)
R_{merge} (%)	6.6 (47.9)	10.2 (55.6)
I/σ (I)	25.4 (2.9)	22.6 (4.1)
Completeness (%)	99.5 (98.5)	99.8 (99.3)
Redundancy	7.3 (6.9)	7.4 (7.1)
Wilson B (Å ²)	44.7	71.4
Refinement statistics		
Resolution (Å)	37.84-2.25 (2.33-2.25)	38.41-2.77 (2.92-2.77)
No. reflections	35446	19437
R_{work}/R_{free} (%)	20.2/23.6 (35.9/35.3)	20.1/25.9 (27.5/34.6)
Number. atoms		
Protein	4967	4848
Ligand/ion	62	62
Water	211	2
Average B-factors (Å ²)		
Protein	53.33	76.13
Ligand/ion	66.53	81.94
Water	42.79	30.00
Rmsds		
Bond lengths (Å)	0.009	0.013
Bond angles (°)	1.337	1.518
Ramachandran plot statistics		
Most favored (%)	91.0	91.9
Additional allowed (%)	8.8	7.5
Generously allowed (%)	0.2	0.7
Disallowed (%)	0.0	0.0

^aAll data sets were collected from a single crystal.

^bValues in the parentheses are for the highest-resolution shell.

ecules (A and B) of BRI1-KD-BIM complex. In both A and B molecules, 16 of the BIM residues (³⁰⁷TMEELQA-AIQAAIAHC³²²) are clearly visible in the electron-density map. The overall structures of BRI1-KD in molecules A and B are similar, and the two structures overlap with one another with a root-mean-square deviation (rmsd) between all C α atoms of 0.47 Å (Supplementary information, Figure S2A and S2B). The structural analyses hereafter are on the basis of molecule A. In the BRI1-KD-BIM complex, the catalytic domain of BRI1 displays the typical protein kinase fold which contains a small N-terminal lobe comprised mainly of β strands, and a large C-terminal lobe that consists mostly of α helices (Figure 2A). The BIM peptide adopts a helical conformation, as confirmed by the well-defined electron density (Supplementary information, Figure S2C), and binds to a groove present between α H and α G in the C-lobe of

BRI1-KD, located distal to the common substrate-binding region. Topologically, the structure of BRI1-KD is similar to that of the active BAK1-KD (PDB ID: 3UIM) [26], with which it shares approximately 40% sequence identity. The overall rmsd of superposition of BRI1-KD with BAK1-KD is 1.04 Å (Figure 2B and Supplementary information, Figure S2E). A sequence alignment based on the structure is presented in Supplementary information, Figure S2D. Differences only occur in particular amino acids, which may determine their distinct function in various signaling pathways (see later).

The activation segment in BRI1-KD (residues Asp1027-Glu1056) is well-ordered and has the conformation typical of all active protein kinases. The structure reveals phosphorylation on the side chains of Thr1039, Ser1042, Ser1044 and Ser1060 in the activation segment of BRI1-KD. The phosphate group of the phosphorylat-

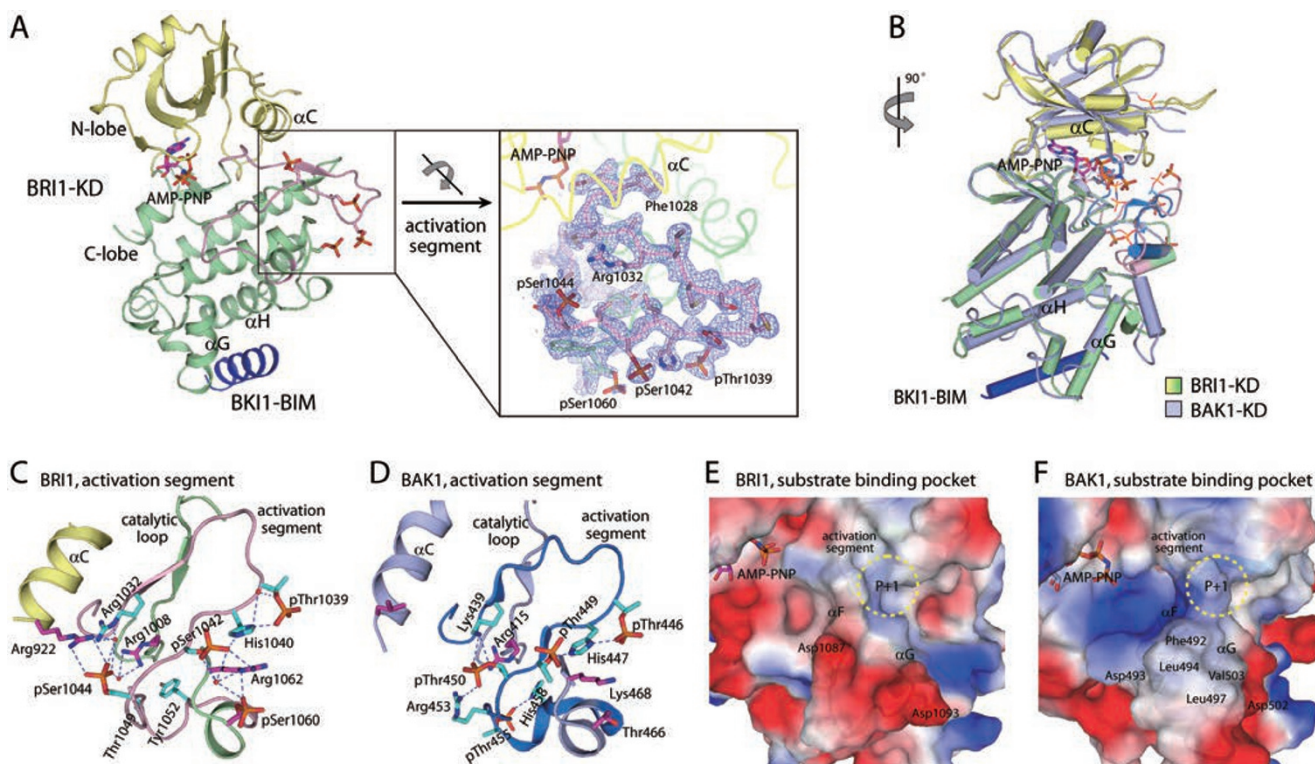


Figure 2 Overall structure of BRI1-KD in complex with BKI1-BIM. **(A)** Ribbon diagram of BRI1-KD-BIM complex. The N-lobe, C-lobe, and the activation segment of BRI1-KD are colored in pale yellow, pale green and pink, respectively, and BKI1-BIM is shown in blue. AMP-PNP and the observed phosphorylated residues are shown as sticks. The coloring scheme of BRI1-KD is the same in the following figures unless indicated. The inset shows the $F_o - F_c$ omit map (contoured at 2.0 σ) for the activation segment of BRI1-KD. **(B)** Comparison of BRI1-KD and BAK1-KD. BAK1-KD is colored in light blue with its activation segment in marine blue, and its phosphorylated residues and bound AMP-PNP are shown as lines. **(C and D)** Interaction networks involving phosphor-Thr/Ser residues in the activation segment of BRI1 **(C)** and BAK1 **(D)**. Residues in the activation segments are highlighted as cyan sticks. Blue dashed lines represent the polar interactions. **(E and F)** Molecular determinants for the substrate specificity of BRI1 **(E)** and BAK1 **(F)**. The classic substrate-binding site are encircled by yellow dashed lines. Key residues potentially determining the substrate specificity of BRI1 and BAK1 are indicated. See also Supplementary information, Figure S2 and Table S1.

ed Ser1044 is positioned to compensate for a positively charged pocket, which is clustered by three arginine residues Arg922, Arg1008 and Arg1032 (Figure 2C and 2D). Residue Ser1044 is the key phosphorylation site corresponding to Thr197 in protein kinase A (PKA), whose phosphorylation is important for promoting the correct orientation and electrostatic environment for the catalytic base Asp1009, as well as the proper conformation of helix α C. Overexpression of BRI1^{S1044A} in *bri1-5* transgenic plants failed to rescue the dwarf phenotype [27], indicating that this residue is important for the activation of BRI1. The other three phosphorylated residues form multiple direct and water-mediated interactions with His1040 and Arg1062. Together, massive networks of hydrophilic interactions stabilize the conformation of the activation segment. The AMP-PNP molecule is well positioned in the ATP-binding site formed by the hinge region, the flexible P-loop in the N-lobe, and the DFG motif in the C-lobe (Supplementary information, Figure S2F). There is no density for Mg²⁺ in the structure although the crystallization condition included sufficient amounts of magnesium. One possible explanation for this is that our crystallization buffer contains 200 mM of citrate which can combine with Mg²⁺ and greatly reduces the concentration of free Mg²⁺ in the crystallization buffer. Similar results have been observed in the crystal structures of other kinases, indicating that the absence of the density for magnesium occurs occasionally for kinase structures [26].

BRI1-KD and BAK1-KD recognize distinct target sequences, and therefore exhibit different substrate specificities. BRI1 is a basophilic serine/threonine kinase that preferentially phosphorylates substrates with basic residues in close proximity to the phosphorylation site, and optimum phosphorylation peptide of BRI1-KD required Arg or Lys residues at P-3, P-4 and P+5 [21]. To provide a structural explanation for substrate selectivity, we compared the surface electrostatic potential of the substrate-binding pockets on BRI1-KD and BAK1-KD (Figure 2E and 2F). The P+1 specificity site of protein kinases, originally named for its role in contacting the P+1 residue in structure of PKA in complex with a peptide inhibitor [28], is formed by a surface loop (residues Thr1045-Glu1056 in BRI1) contiguous with the activation loop (residues Ser1035-Ser1044 in BRI1). The P+1 site of BRI1-KD is encircled by acidic side chains of Asp1087, Asp1093 and probably pSer1060, which is consistent with the singular preference of BRI1 for positively charged residues around the phosphorylation site on the substrate (Figure 2E). In contrast, the corresponding region of BAK1-KD consists of several hydrophobic residues (Phe492, Leu494, Leu497 and Val503), while

the acidic Asp493 and Asp502 point in the opposite directions to that of the corresponding Asp1087 and Asp1093 in BRI1-KD (Figure 2F). These subtle structural differences between BRI1-KD and BAK1-KD may determine their distinct functions in various signaling pathways by recognizing different substrates.

Docking interactions between BIM peptide and BRI1 KD

Complex formation between BIM and BRI1-KD buries a surface area of $\sim 638 \text{ \AA}^2$. The traceable region of the BIM peptide folds into a four-turn α -helix and is embraced by α G and α H from BRI1-KD (Figure 3A). The BIM peptide lies in a hydrophobic groove formed by residues Val1100, Lys1101, Ala1104 and Ile1108 from helix G and its succeeding loop at one side and residues Ile1125, Leu1128, Gln1129, Leu1131, Val1135 and Leu1138 from helix α H at the opposite side. Six hydrophobic residues (Met308, Leu311, Ala314, Ile315, Ala318, Ile319) and the methylene group of Gln312 from the BIM peptide are involved in the hydrophobic interaction with BRI1-KD. In addition, Lys1132 from helix α H of BRI1-KD forms a hydrogen bond with Gln316 in the BIM peptide of BKI1.

To assess the importance of the aforementioned interactions, we generated a series of point mutations on the BKI1-Cter and examined their effect on the Ser270/Ser274 phosphorylation of BKI1-Cter catalyzed by BRI1-KD (Figure 3B). When the key hydrophobic residues on the BIM sequence were individually replaced by charged residues or Ala (BKI1-Cter^{M308D}, BKI1-Cter^{L311D}, BKI1-Cter^{I315D}, BKI1-Cter^{A318D} and BKI1-Cter^{I319A}), their transphosphorylation rates by BRI1-KD showed a 2-6-fold decrease. The triple mutations (BKI1-Cter^{M308D/A314D/A318D} and BKI1-Cter^{L311D/I315D/I319A}) resulted in a further decrease in BRI1-KD's transphosphorylation activity towards BKI1-Cter, indicating a negatively synergistic effect in BRI1-KD-BKI1-Cter interaction. As the negative controls, we also made a mutant, BKI1-Cter^{Q312A}, and assessed its effect on phosphorylation rate. As expected, no significant effect was observed for this mutant. Interestingly, the Q316E mutation yielded a modest but marked increase in its phosphorylation rate by BRI1-KD, probably because Glu316 in this mutant formed a salt bridge with Lys1132 of BRI1-KD which enhances the BRI1/BKI1 interaction. BKI1 represents a class of membrane associated kinase regulators (MAKRs) with 7 members in *Arabidopsis*, which contain both the N-terminal and C-terminal linear motifs. Jaillais *et al.* [15] have shown that some members of MAKR family (MAKR3 and MAK4) could not interact with BRI1-KD *in vitro*, and their overexpression did not result in a BR-related phenotype. Secondary structure predictions

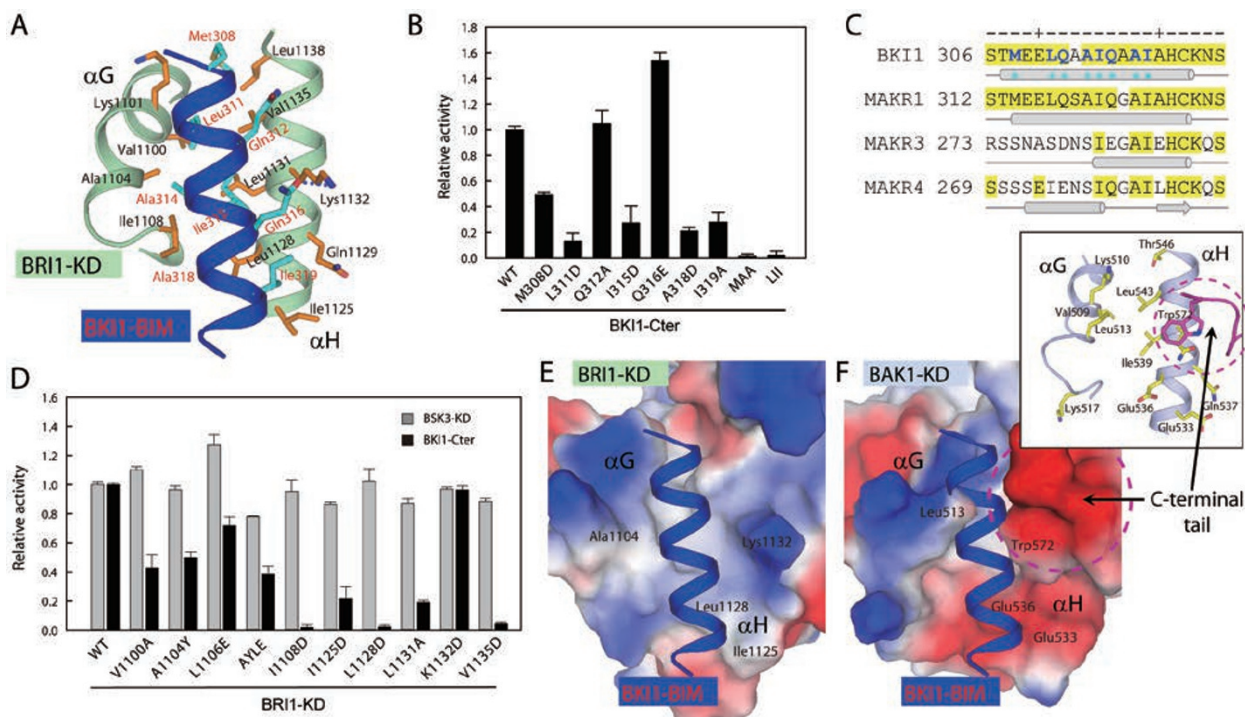


Figure 3 BKI1-BIM specifically recognizes BRI1-KD. **(A)** Interactions between BKI1-BIM and BRI1-KD. Residues from BKI1-BIM and BRI1-KD are highlighted as cyan and orange sticks, respectively. **(B)** Effects of BKI1 mutants on the phosphorylation efficiencies of BRI1-KD towards BKI1-Cter fragment (mean \pm SEM, $n = 3$). **(C)** Sequence alignment of the BIM regions from BKI1 and MAKRs. The UniProt identifiers for BKI1, MAKR1, MAKR3, and MAKR4 are Q9FMZ0, Q3E936, Q9ZUS8 and O80624, respectively. The key residues required for BRI1-BKI1 recognition are indicated by cyan triangles. The observed and predicted secondary structure assignments are shown below each sequence. **(D)** Effects of BRI1-KD mutants on its phosphorylation efficiencies towards substrates BKI1-Cter and BSK3-KD (mean \pm SEM, $n = 3$). **(E and F)** Comparison of the docking grooves for BKI1-BIM on BRI1-KD **(E)** and BAK1-KD **(F)**. The BRI1-KD and BAK1-KD are shown in surface representation colored according to electrostatic potential (positive, blue; negative, red), and the BKI1-BIM peptide is highlighted as blue ribbon. The hindrance of Trp572 from the C-terminal tail of BAK1-KD for BKI1-BIM binding is shown in the inset of **F**.

using the procedure of Rost and Sander [29] delineated residues 273–281 in MAKR3 and 279–288 in MAKR4 as a region of highest non-helix probability (Figure 3C), which indicates that integrity of the α -helical structure is also required for BKI1’s inhibitory function.

Next, we systematically substituted key residues in the BRI1 interface and determined the contribution of individual residue to the BRI1-KD-catalyzed reactions. As shown in Figure 3D, all the mutants had similar catalytic activity towards BSK3-KD substrate. In contrast, a dramatic reduction of BRI1-KD activity for BKI1-Cter in the range of 25–50-fold was observed upon replacement of three crucial hydrophobic residues with an Asp (BRI1-KD^{I1108D}, BRI1-KD^{L1128D} and BRI1-KD^{V1135D}). Unexpectedly, mutating Lys1132 to charged Asp had little effect on enzyme activity, suggesting that this basic residue is not important for BKI1 targeting compared to the hydrophobic residues. BKI1-Cter specifically binds to BRI1-KD but not the KD of the coreceptor BAK1 *in*

vitro [14, 15]. We aligned the C-lobes of the KDs from the crystal structures of BAK1-KD and BRI1-KD-BIM complex as shown in Figure 3E and 3F. One of the few differences between BRI1-KD and BAK1-KD in the BIM-binding region is the presence of Ile1125/Leu1128 in BRI1-KD, which correspond to Glu533/Glu536 in BAK1-KD. The substitution of the two hydrophobic residues with negatively charged residues may completely disrupt the hydrophobic environment required for BKI1 binding on BAK1. In addition, the small residue Ala1104 in BRI1 is replaced by Leu513 with large side chain in BAK1. When we modeled BIM peptide onto the BAK1 KD by using the structure of BRI1-KD-BIM complex as a guide, Leu513 in BAK1 collided with the middle portion of the BIM helix. The Cter tail of BAK1-KD (residues 566–573) is ordered and positioned on the top of helix α G, where the bulky indole ring of Trp572 would sterically clash with the hypothetically bound BIM peptide. Together, the acidic patch and two bulky side chains in

the BAK1-KD would impede the binding of the BIM peptide.

BRI1-BKII interaction is necessary to inhibit BRI1 signaling in planta

A key step in BRI1 activation is the dissociation of BKII from the plasma membrane (PM), and the phosphorylation of BKII by BRI1 plays critical roles in this process [14, 15, 19]. To explore the biological relevance of the interaction of BKII with BRI1, transgenic plants expressing the interface-mutated BKII-GFP were generated. We found that none of the transgenic seedlings showed a phenotype similar to *BKII* overexpression lines. As shown in Figure 4A, overexpression of the wild-type *BKII* resulted in dwarf plants that had a smaller rosette than the untransformed plants. In comparison with the wild-type *BKII*, *BKII*^{L311D} overexpression resulted in a semidwarf phenotype, but overexpression of the other six mutants defective for interaction with BRI1 failed to produce a dwarf phenotype, despite these proteins being expressed at levels similar to wild-type *BKII* (Figure 4B). Interestingly, *BKII*^{Q316E} overexpression resulted in severe dwarf plants that resembled the

phenotype of transgenic plants with overexpression of myristoylated BKII [14]. Indeed, Q316E substitution in BKII-Cter increased its interaction with BRI1-KD ($K_m = 2.34 \pm 0.44 \mu\text{M}$ for BKII-Cter^{Q316E} vs $K_m = 4.28 \pm 0.73 \mu\text{M}$ for wild-type BKII-Cter).

We also examined the subcellular distribution of BKII mutant forms in plants (Figure 4C). Like wild-type BKII-GFP, BKII^{Q316E}-GFP was localized to both the plasma membrane (PM) and the cytosol in root-tip cells, and BRs induced the dissociation of this mutant from the PM, as indicated by the diminished discrete fluorescent signal on the cell surface. In contrast, BRs did not cause complete dissociation of other BKII-GFP mutants from the PM. The localizations of the BKII-GFP mutants are consistent with their phosphorylation efficiencies by BRI1, further confirming that release of BKII from the PM is triggered by multiple phosphorylation. Note that the M308D mutant appears to have some internalized BKII, more than the partial loss-of-function mutant L311D, suggesting that the phenotype of the transgenic plant is not always correlated with BKII localization. Similar result has been reported by Jaillais *et al.* [15]. They found that although the BKII^{C322A} transgenic plant

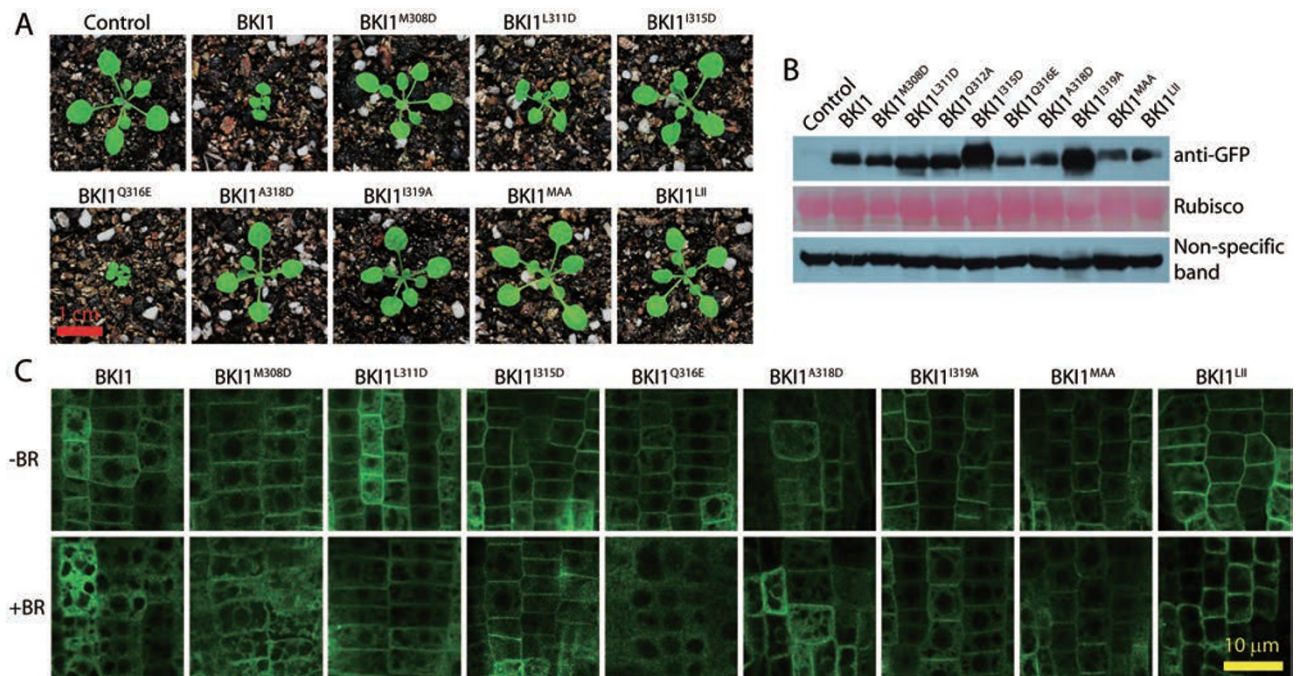


Figure 4 BKII mutants defective in BRI1 interaction lose their inhibitory activity in plants. **(A)** Phenotypes of plants expressing empty vector and various *BKII*-GFP in the Columbia (Col-0) *Arabidopsis*. Scale bar, 1 cm. For each construct, 30 transgenic lines with comparable levels of BKII expression were selected for further phenotypic analysis. The rosette radius of the plants were measured and the phenotypes in each group are consistent (Standard deviations for all data sets in each group were less than 10% of the mean). **(B)** Protein levels of various BKII-GFP in transgenic plants. Ponceau-S-stained Rubisco large subunit served as a loading control. **(C)** Localization and membrane dissociation upon 1 μM epi-BL treatment of various BKII-GFP in root tip cells of 4-day-old seedlings. Scale bar 10 μm .

showed normal growth as the Col-0, it had abundant internalized BKI1^{C322A}.

BKI1 binding to BRI1 inhibits the reciprocal transphosphorylation of BRI1 and BAK1

The PM-localized BRI1 is quiescent in the absence of steroids, largely due to a negative regulator, BKI1 [14, 15, 19]. BKI1 could control BRI1 activity by two possible mechanisms: (1) directly inhibit the intrinsic kinase activity of the activated BRI1 by blocking its active site, or (2) inhibit the activation process of the inactive BRI1. To distinguish these mechanisms, we first measured the

binding affinities of the unphosphorylated and phosphorylated BKI1-Cter (pBK11-Cter) with BRI1-CD by isothermal titration calorimetry (ITC; Figure 5A). The phosphorylated BKI1 has lower affinity than dephosphorylated BKI1 to BRI1 KD ($K_d = 3.95 \mu\text{M}$ vs $1.61 \mu\text{M}$), which is plausible as phosphorylated BKI1 leaves PM and BRI1 in the presence of BRs. Furthermore, Wang *et al.* [16] have shown that the 14-3-3 proteins can interact with the phosphorylated BKI1 through a motif that contains the two phosphorylation sites to release inhibition of BRI1 by BKI1.

Because pBK11-Cter is a product of the BRI1-CD-cat-

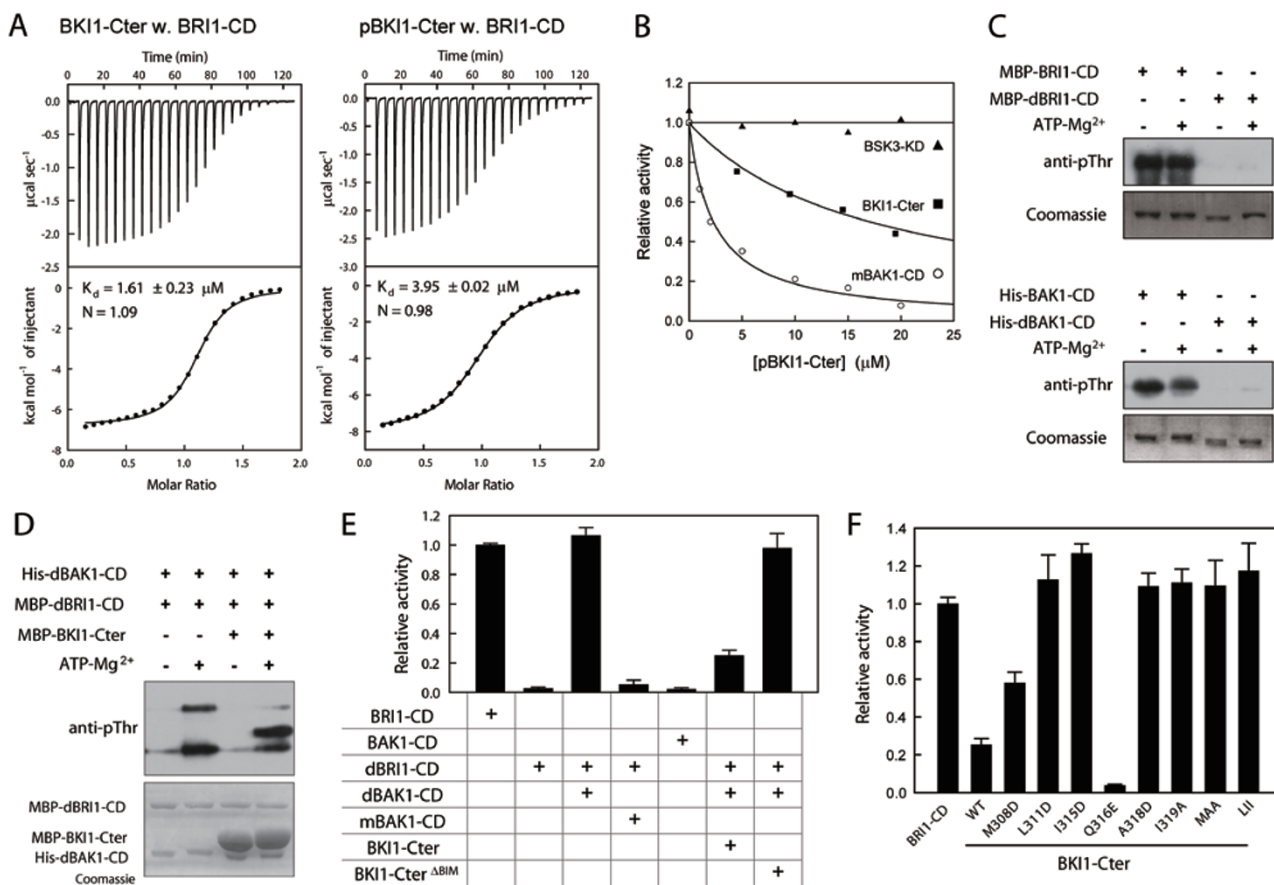


Figure 5 BKI1 inhibits transphosphorylation of BRI1 and BAK1. **(A)** Thermodynamic analysis of the interaction between BKI1-Cter or pBK11-Cter and BRI1-CD by ITC. The top panels show heat release during titration, and the bottom panels show the corresponding integrated heat of binding. **(B)** Effects of pBK11-Cter on the BRI1-catalyzed phosphorylation reactions. The relative activity of BRI1-CD towards substrates BSK3-KD, BKI1-Cter and mBAK1-CD were measured in the presence of pBK11-Cter. Reaction was initiated by adding 50 nM BRI1-CD to the kinase reaction buffer containing 5 µM substrate (BSK3-KD, BKI1-Cter or mBAK1-CD) in the presence of different concentrations of pBK11-Cter. The initial velocities of BSK3-KD, BKI1-Cter and mBAK1-CD phosphorylation were determined. **(C)** Autophosphorylation of various BRI1-CD and BAK1-CD proteins analyzed by western blotting. BRI1-CD or BAK1-CD purified from *E. coli* was incubated with GST-PP2Cα *in vitro* to produce the dephosphorylated BRI1-CD (dBRI1-CD) and the dephosphorylated BAK1-CD (dBAK1-CD). The dephosphorylated proteins were further purified to remove the phosphatase. **(D and E)** Effects of BKI1-Cter on transphosphorylation of dephosphorylated BRI1-CD and BAK1-CD analyzed by western blotting **(D)** and enzyme kinetic assays **(E)**. **(F)** Inhibition of various BKI1 mutations on BRI1-BAK1 transphosphorylation (mean ± SEM, *n* = 3).

alyzed reaction and has high affinity for BRI1-CD, we then evaluated the ability of pBK11-Cter to inhibit the BRI1-catalyzed reactions. As shown in Figure 5B, pBK11-Cter efficiently inhibits the BRI1-CD-catalyzed phosphorylation of BK11-Cter and mBAK1-CD, but has no effect on the BSK3-KD phosphorylation. This is consistent with the result obtained by *in vitro* GST pull-down assay that BK11 directly inhibited the interaction between the KDs of BRI1 and BAK1 [14]. Taken together, these observations indicate that (1) BK11-Cter recognizes BRI1-CD independent of its own phosphorylation status; (2) the binding of BK11-Cter and BSK3-KD is not mutually exclusive, suggesting that BK11 and BSK3 interact with different regions of BRI1.

Next, we performed two complementary experiments to further examine whether BK11 affects the activation process of the inactive BRI1. Bacterially expressed kinases are often associated with high levels of autophosphorylation that is caused by the rapid overexpression of the kinases in a strong expression system, leading to phosphorylation of proteins before or during protein folding. Therefore, phosphorylation events during expression may occur at non-physiological sites that are not accessible as substrate sites in folded proteins [30]. To investigate whether the observed hyper-phosphorylation of BRI1-CD and BAK1-CD is non-physiological or not, *in vitro* autophosphorylation experiments were performed on BRI1-CD and BAK1-CD, starting with non-phosphorylated proteins. To produce the dephosphorylated BRI1-CD and the dephosphorylated BAK1-CD, BRI1-CD or BAK1-CD expressed and purified from *E. coli* was incubated with GST-PP2Ca at a 20:1 ratio for 1 h at 25 °C in the buffer containing 10 mM MgCl₂ and 100 mM NaCl. After incubation, GST-PP2Ca was removed by GST columns. The dephosphorylated proteins were further purified by gel filtration chromatography. The dephosphorylated form of BRI1-CD (2 μM) was incubated with ATP-Mg²⁺ (1 mM ATP and 10 mM Mg²⁺) either in isolation or together with the dephosphorylated form of BAK1-CD (2 μM) for 1 h, followed by SDS-PAGE. As shown in Figure 5C, neither dBRI1-CD nor dBAK1-CD alone is able to undergo ATP-initiated autophosphorylation. However, when the dephosphorylated forms of BRI1-CD and BAK1-CD were mixed together in the presence of ATP-Mg²⁺, both of them were phosphorylated. The presence of the BK11-Cter greatly decelerated the transphosphorylation of BRI1-CD and BAK1-CD (Figure 5D). These results seem to be inconsistent with those obtained by other groups. Oh *et al.* [21] first proposed that bacterially expressed BRI1 alone can undergo an intramolecular autophosphorylation *in vitro*, while Wang *et al.* [13] proposed that BRI1 autophos-

phorylation was intermolecular [21]. One of the primary differences between our and other studies is that bacterially expressed BRI1 used in their studies were active and phosphorylated at multiple sites, while we used dephosphorylated BRI1 for autophosphorylation assay. Thus, our studies demonstrated that the initiation of BRI1 autophosphorylation requires the presence of BAK1.

We also performed *in vitro* kinase assays to examine the BRI1-CD activity with BSK3-KD substrate during the reciprocal phosphorylation of these two KDs. Under the same experimental conditions, the activation of BRI1-CD is well correlated with its phosphorylation level (Figure 5E). In addition, all the BK11 mutants defective for interaction with BRI1 impaired inhibition for the BAK1-mediated BRI1 activation (Figure 5F). These results indicated that (1) the initial activation of either receptor kinase after hetero-dimerization is through the reciprocal transphosphorylation rather than *cis*-autophosphorylation; (2) binding of BK11 to BRI1-CD blocks the interaction between the BAK1 and BRI1 KDs [14], and prevents the reciprocal transphosphorylation.

Discussion

In this study, we have determined the structure of the BRI1 KD in complex with a docking peptide derived from BK11. While the current manuscript was under review, crystal structures of BRI1 KD in isolation and in complex with adenine nucleotides (PDB codes: 4OA2, 4OA6, 4OA9, 4OAB and 4OAC) were reported [31]. These five structures are almost identical and readily superimposable to our BRI1 KD with overall rmsd of < 0.5 Å over ~220 Ca atoms (Figure 6A and Supplementary information, Figure S3A). In our structure, the non-hydrolyzable ATP analogue AMP-PNP adopts a catalytically incompetent configuration, similar to that observed in the BRI1-KD-ATP complex structure (4OAB; Supplementary information, Figure S3B and S3C). The γ-phosphate rotated outwards away from the catalytic Asp1009, different from its active conformation in the BRI1-KD/AMP-PNP/Mn²⁺ complex structure (4OA9). On close inspection, however, we found significant discrepancies in the region of residues 1087-1124, including helices αG, αG', αG'' and their flanking loops (termed the αG region) while superimposing our BRI1-KD/AMP-PNP/BIM complex to their high-resolution representative (4OAC) (Figure 6B and 6C). The amino acid sequence of this αG region in their BRI1-KD structure is off-registered by up to 7 amino acids relative to our structure. As a consequence, key residues for interaction with the BK11-BIM peptide, such as Val1100, Lys1101, Ala1104 and Ile1108, are “frame shift” by approximately

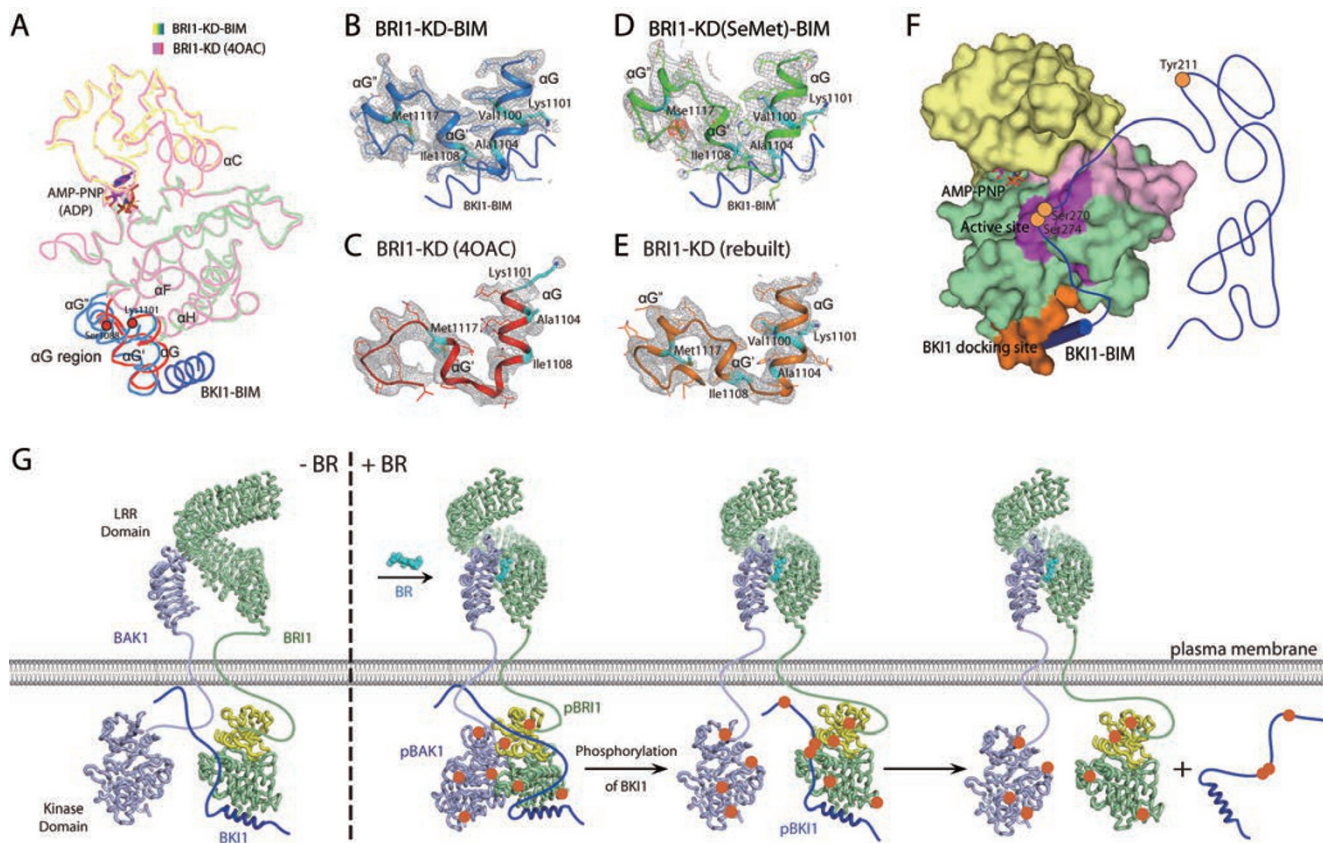


Figure 6 Model for early events of BR signaling. **(A)** Superposition of our BRI1-KD-BIM complex with the reported BRI1-KD structure (4OAC). The α G regions of residues 1 087-1 124 in our and reported structures are highlighted in marine blue and red, respectively. **(B-E)** Close-up views of the α G regions in our native **(B)** and SeMet-labeled **(D)** BRI1-KD-BIM complexes and in the reported **(C)** and rebuilt **(E)** BRI1-KD structures. The $2F_o - F_c$ electron density maps (contoured at 0.8σ) are shown in gray mesh. In panel **D**, the anomalous difference Fourier map around the seleno-methionine Mse1117 (contoured at 3.0σ) is shown in red mesh. Key residues for BKI1 interaction are highlighted in cyan sticks. **(F)** Interaction model of BRI1 and BKI1. BRI1-KD is shown in surface representation, and the phosphorylation sites on BKI1 are shown as orange dots. **(G)** Working model for the regulation of BR signaling receptor complexes. Without BR, BKI1 competitively inhibits the transphosphorylation of BAK1 and BRI1. BR-induced heterodimerization of BRI1 and BAK1 LRR domains brings their cytoplasmic regions in the right orientation to remove BKI1 inhibition and trigger transphosphorylation of two kinase domains. The activated BRI1 then phosphorylates BKI1, which accelerates its membrane dissociation. The BRI1-BAK1 complex is fully activated. Notably, the initiation of BR signaling may also involve a small amount of preassembled BRI1/BAK1 heterodimers or BRI1 homodimers [39, 40]. See also Supplementary information, Figure S3 and Table S2.

two helix turns. In their reported structure, the preceding loop of helix α G (1 089-1 100) was disordered (Figure 6A). Although the electron density for the rest of the α G region is largely continuous over the main chain, there is insufficient density to justify building side-chains (Figure 6E and Supplementary information, Table S2). One obvious question is whether the observed differences in the α G region are modulated by BKI1 binding. To address this issue, we first used selenium anomalous signals to help validate the atomic model of our BRI1-KD-BIM complex (Supplementary information, Figure S3D and Table 1). Notably, residue Met1117 is indubita-

bly located on helix α G'' (Figure 6D), indicating that the assignment of the α G region in our structure is reliable. In addition, BRI1 and BAK1 share approximately 70% sequence similarity in the KD and 60% similarity in the α G region, making direct comparison between these two proteins relevant. There are two reported structures of the BAK1-KD, one in isolation (3UIM) and the other in complex with AvrPtoB (3TL8) [26, 32]. The α G region of BAK1-KD mediates the tight interaction with the effector protein AvrProB. Structural analyses and sequence alignment reveal that both BAK1-KDs adopt a conformation in the α G region very similar to that in our BRI1-

KD-BIM structure (Supplementary information, Figure S3A). Therefore, we believe that the conformation of the α G region in BRI1-KD is independent of BKI1 binding.

Structural and mutagenesis studies showed that the α G region of BRI1 is required for recognition of the BKI1-BIM peptide, and shed light on the importance of this interface for BKI1-mediated inhibition. BKI1 has three phosphorylation sites that are 30-100 residues away from the BIM sequence [15, 19]. These sites are predicted to reside in the disordered polypeptide regions, suggesting that these regions are dynamic when bound to BRI1 and that interactions at the BIM motif tether the substrate to BRI1 allowing the phosphosite(s) to encounter the catalytic site and be phosphorylated (Figure 6F). Several mechanical events have been shown to contribute to the ligand-induced activation of BRI1. These include ligand-induced conformational change in a preformed homodimer or receptor homodimerization and autophosphorylation of BRI1 KD, phosphorylation and disassociation of BKI1 from the PM, and association and transphosphorylation with the BAK1 coreceptor kinase [17, 33]. In order to determine the exact sequence and causal relationship of these events, we investigated the detailed process of BRI1/BAK1 activation with dephosphorylated BRI1-CD and BAK1-CD. Current and previous studies provide several lines of evidence to strongly support that phosphorylation of BRI1 by BAK1 is indispensable to the early events of the BR signaling pathway. The evidence for this may be summarized as follows:

(1) Bacterially expressed KDs of BRI1 and BAK1 are phosphorylated at multiple sites when purified from *E. coli*, while the corresponding kinase-dead mutants, mBRI1-CD and mBAK1-CD, were completely unphosphorylated (Figure 1). BRI1-CD, but not BAK1-CD, can phosphorylate mBAK1. In contrast, BAK1-CD, but not BRI1-CD, can phosphorylate mBRI1-CD.

(2) Although bacterially expressed BRI1-CD and BAK1-CD are catalytically active, neither dephosphorylated BRI1-CD nor BAK1-CD is able to undergo activating autophosphorylation when incubated alone with ATP-Mg²⁺ *in vitro*. In contrast, both of them can be phosphorylated and activated when they were incubated together in the presence of ATP-Mg²⁺ (Figure 5). These results suggest that the reciprocal transphosphorylation on the cytoplasmic domains of BRI1 and BAK1 in the complex is a prerequisite for the full activation of the receptor kinase.

(3) In yeast, BRI1-BAK1 interaction activates both receptor kinases. Little phosphorylation was detected for BRI1 when it was expressed alone, however, both BRI1 and BAK1 were phosphorylated when they were coexpressed in yeast cells. Yet, no phosphorylation on either

receptor kinase was detected when one of the receptor kinases was mutated. These results imply that the full activation of either receptor kinase after heterodimerization is through transphosphorylation rather than *cis*-phosphorylation [5].

(4) BAK1 is a conserved member of the SERK subfamily of LRR-RLKs [34]. Genetic and biochemical studies showed that four out of five members of the SERK subfamily (SERK1 to SERK4) may play functionally redundant roles in BR signaling. Detailed analysis indicated that, without BAK1 and its functionally redundant proteins, BR signaling is completely disrupted, largely because BRI1 has lost its ability to activate downstream components [35]. This study provided clear genetic and biochemical evidence that BRI1 cannot initiate BR signaling without BAK1 and its homologues. The structure of BRI1-BL-SERK1 complex revealed that the BRI1-LRR and SERK1-LRR form heterodimer in a BR-dependent manner, suggesting that BRs act as “molecular glues” for receptor-coreceptor interaction [36].

Based on the results presented in this study and the previous knowledge on BR signaling, we propose the following model for the activation of a membrane receptor upon BR binding (Figure 6G). In the absence of steroid ligand, the BR-binding site (a region that includes ~70 amino acid ‘island’ domain) in the BRI1 extracellular domain is largely disordered [37, 38], and the cytoplasmic KD of BRI1 associates tightly with BKI1. The BRI1-BKI1 complex limits the interaction of BRI1 with BAK1 KD and prevents the activation of BRI1. Hormone binding induces a conformational rearrangement and fixing of the island domain. The steroid ligand and the now fully ordered island domain together form a docking platform that allows for the steroid-induced binding of the small coreceptor kinase BAK1 [33, 36]. Physical interaction between the BRI1 and BAK1 extracellular domains brings their cytoplasmic KDs closer together, and makes them competent to transphosphorylate each other. BAK1 has a low basal kinase activity that is responsible for the BRI1 phosphorylation upon heterodimerization. The major function of BRI1-BAK1 complex is to increase the “effective concentration” of the phosphorylation sites of the BRI1 in the vicinity of the active site of BAK1, and hence enhances the catalytic efficiency for BRI1 phosphorylation. The BAK1-activated BRI1 KD is able to rapidly phosphorylate multiple substrates, such as BAK1, BKI1 and BSK3, for activating downstream signaling responses [4, 5, 14, 18]. The phosphorylated BKI1 dissociates from PM and BRI1, and then binds to 14-3-3 proteins in cytosol to free BES1/BZR1. The released BES1/BZR1 accumulates in the nucleus, leading to BR-responsive gene expression [15, 19]. The BRI1-acti-

vated BAK1, in turn, transphosphorylates BRI1 rapidly, leading to enhanced BR signaling and increased plant growth.

Recently, Bücherl *et al.* [39] investigated the spatial correlation between BRI1 and BAK1 in the PM of live *Arabidopsis* root epidermal cells using comparative colocalization analysis and fluorescence lifetime imaging microscopy. They showed that approximately 7% of the BRI1 PM pool constitutively heterodimerizes with BAK1 in live root cells and activation of BR signaling by exogenous ligand application resulted in an about 50% increase of receptor heterodimerization in the PM. Our model can be extended to include the contribution of the BR-independent BRI1-BAK1 heterodimerization equilibrium (Supplementary information, Figure S3E). This model may also expand beyond BRI1/BAK1 pair of receptors due to the multi-functions of BAK1 and its homologues. A flagellin-induced FLS2-BAK1 complex activates and initiates plant defense signaling probably in a similar manner to BRI1-BAK1. Moreover, BAK1 and its homologues also function as coreceptors for various ligand-binding LRR-RLKs, such as BIR1 and PEPR1, to regulate cell death and immunity-related signaling pathways [34].

Materials and Methods

Constructs, mutagenesis and protein purification

Fragments of BRI1, BAK1, BKI1 and BSK3 were overexpressed in BL21(DE3) cells as His-tagged and/or MBP-fusion proteins. Mutations were generated by an overlap PCR procedure. All proteins were purified over Ni-NTA or amylose resin, ion-exchange and gel filtration chromatography.

Kinase assays for BRI1

The kinase activity of BRI1 was determined with BSK3-KD or BKI1-Cter as substrates using a coupled spectrophotometric assay. The kinetic parameters were obtained by fitting the experimental data to the Michaelis-Menten equation.

Crystallography

Crystals of the native or SeMet-labeled BRI1-KD-BIM were grown by mixing the protein complex with an equal volume of reservoir solution containing 0.1 M HEPES (pH 7.5), 20% PEG 3350 and 0.2 M Li Citrate. The diffraction data sets were collected at beamline BL17U at the Shanghai Synchrotron Radiation Facility, and the structure was solved by molecular replacement. The data processing and refinement statistics were summarized in Table 1.

Confocal imaging of transgenic plants

Floral dip method was used to generate transgenic lines of wild-type and mutated BKI1-GFP in the Columbia background. Confocal microscopy images of various BKI1-GFP in T₂ seedlings were taken with a Leica SP8 microscope using 488 nm excitation. For exogenous BR treatment, the root tips were first observed without and then treated with 1 μ M epi-brassinolide for 10 min on

slides.

More methods and details were described in Supplementary information, Data S1.

Accession numbers

The atomic coordinates of the native and SeMet-labeled BRI1-KD-BIM complex have been deposited in the Protein Data Bank under accession codes 4OH4 and 4Q5J, respectively.

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(Supplementary information is linked to the online version of the paper on the *Cell Research* website.)