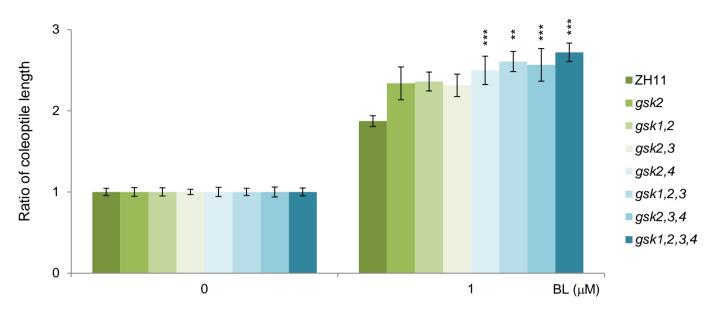
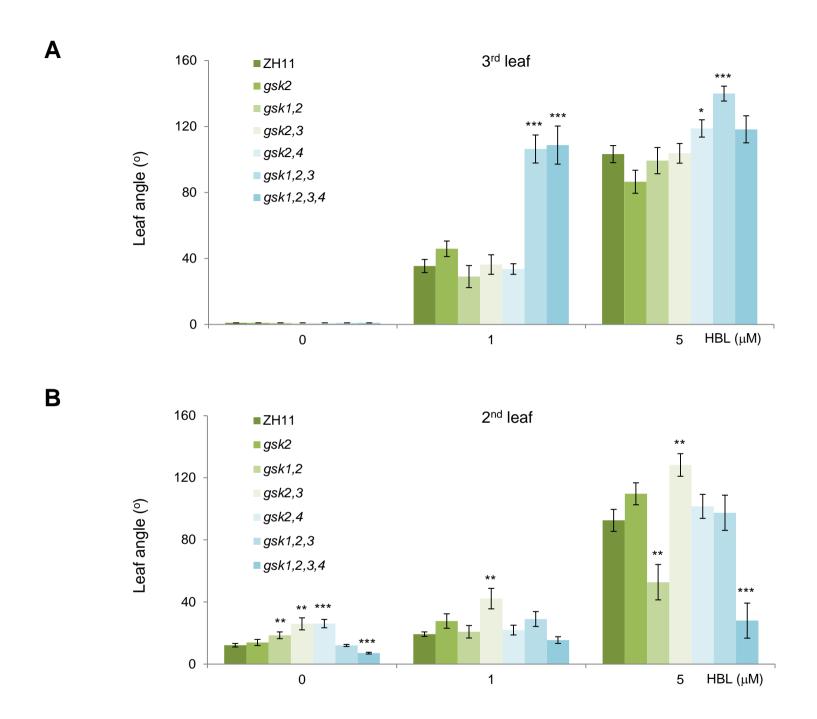


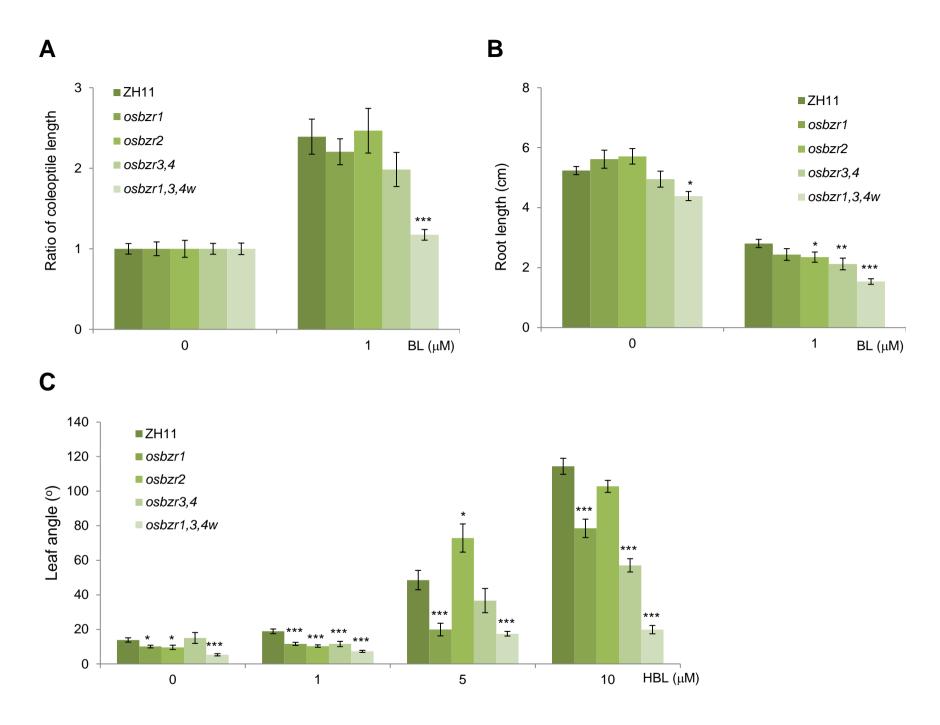
Supplemental Figure S1. Tiller number and root length of *GSK***-related mutants.** (A) Tiller number was counted at the harvesting stage. Bars indicate SD (n = 10). *P < 0.05 generated by t-test. (B) Root length was measured after ten-day growth on agar media. Bars indicate standard error of mean values (SEM, n = 14 - 30). ***P < 0.001 generated by t-test.



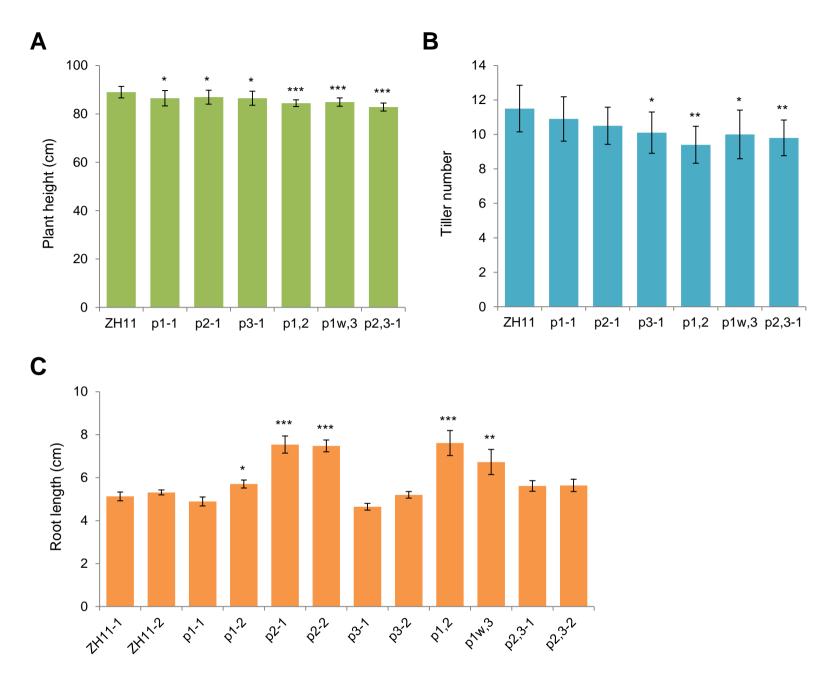
Supplemental Figure S2. BR sensitivity of *GSK*-related mutants evaluated by coleoptile elongation test. Coleoptile length was measured after ten-day growth on agar media with or without BL, and the ratios of BL-treated plants on those untreated were shown. Bars indicate SEM (n = 11 - 30). **P < 0.01 and ***P < 0.001 generated by t-test.



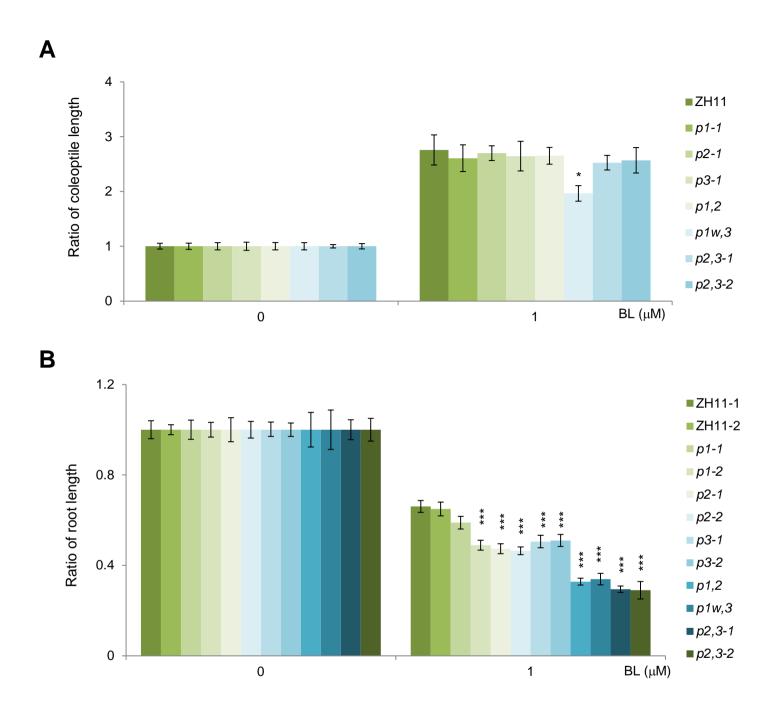
Supplemental Figure S3. BR sensitivity of GSK-related mutants evaluated by lamina bending tests of the 2^{nd} and 3^{rd} leaves. HBL (28-homobrassinolide) was used as BR for the analysis. HBL was applied at 3 days after germination when the 2^{nd} leaves were emerging and leaf angles were measured after additional 4-d-growth. Note that at the time measuring the angles of the 3^{rd} leaves, those without HBL treatment have not inclined thus were counted as zero. Bars indicate SEM (n = 15 - 24). *P < 0.05, **P < 0.01 and ***P < 0.001 generated by t-test.



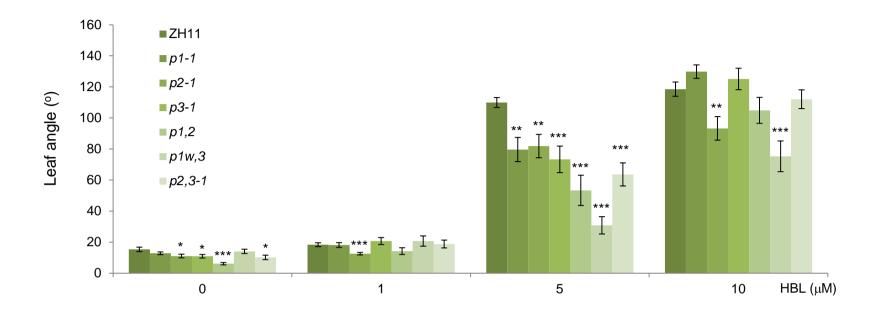
Supplemental Figure S4. BR sensitivity tests of *OsBZR***-related mutants.** (A) Coleoptile length was measured after ten-day growth on agar media, and the ratios of BL-treated plants on those untreated were shown. Bars indicate SEM (n = 8 - 21). ***P < 0.001 generated by *t*-test. (B) Root length was measured after ten-day growth on agar media. Bars indicate SEM (n = 8 - 25). *P < 0.05, **P < 0.01 and ***P < 0.001 generated by *t*-test. (C) HBL was applied at 3 days after germination when the 2nd leaves were emerging and angles (the 2nd leaves) were measured after additional 4-d-growth. Bars indicate SEM (n = 20 - 26). *P < 0.05, **P < 0.01 and ***P < 0.001 generated by *t*-test.



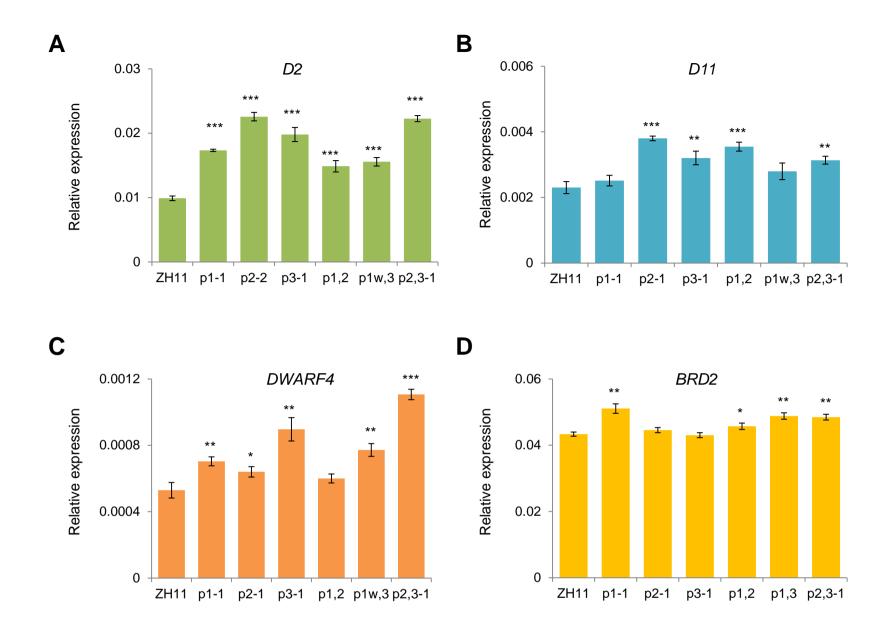
Supplemental Figure S5. Plant height, tiller number, and root length of *PPKL*-related mutants. (A) Plant height was measured at the harvesting stage. Bars indicate SD (n = 16). *P < 0.05 and ***P < 0.001 generated by *t*-test. (B) Tiller number was counted at the harvesting stage. Bars indicate SD (n = 10). *P < 0.05 and **P < 0.01 generated by *t*-test. (C) Root length was measured after ten-day growth on agar media. A replicate of ZH11 was involved for the analysis. Bars indicate SEM (n = 19 - 22). *P < 0.05, **P < 0.01 and ***P < 0.001 generated by *t*-test.



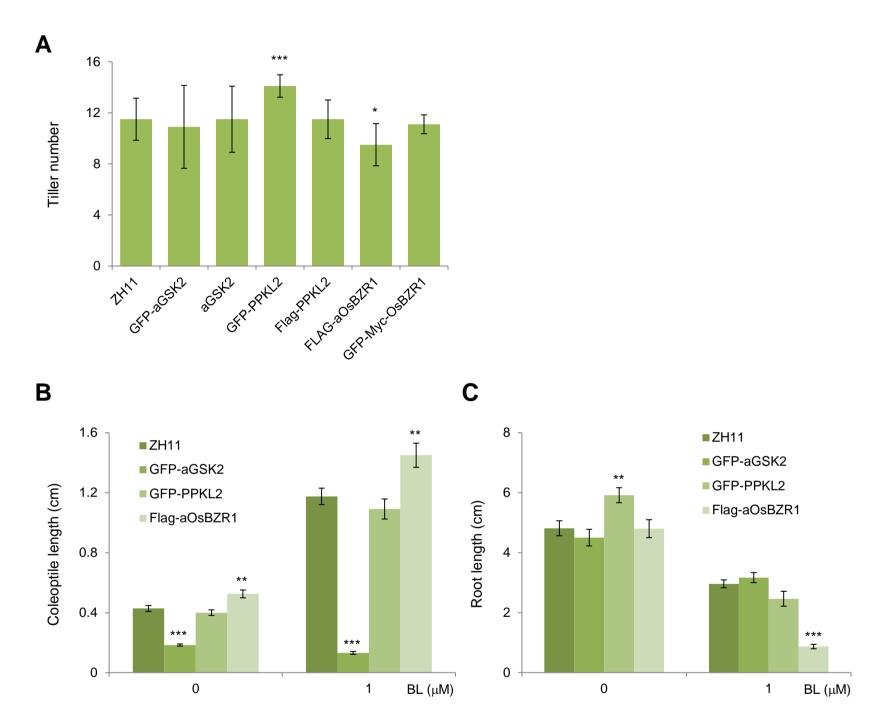
Supplemental Figure S6. BR sensitivity of *PPKL*-related mutants evaluated by coleoptile elongation test and root inhibition test. Coleoptile length and root length were measured after ten-day growth on agar media, and the ratios of BL-treated plants on those untreated were shown. Bars indicate SEM (n = 15 - 26). *P < 0.05 and ***P < 0.001 generated by t-test.



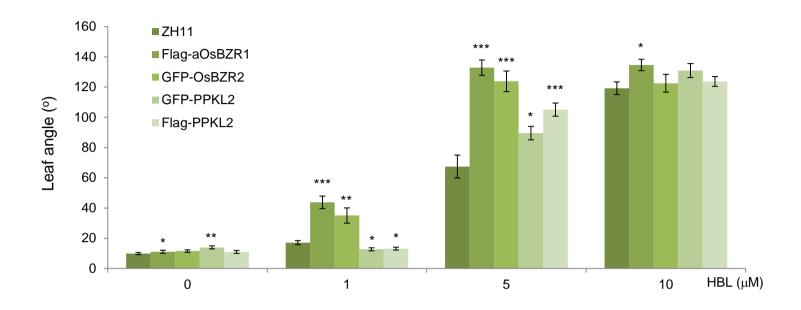
Supplemental Figure S7. BR sensitivity of *PPKL*-related mutants evaluated by lamina bending test. HBL was applied at 3 days after germination when the 2^{nd} leaves were emerging and angles (the 2^{nd} leaves) were measured after additional 4-d-growth. Bars indicate SEM (n = 16 - 31). *P < 0.05, **P < 0.01 and ***P < 0.001 generated by t-test.



Supplemental Figure S8. Expression of BR-synthetic genes in the shoots of one-week-old seedlings of different plants. Bars indicate SD (n = 3). *P < 0.05, **P < 0.01 and ***P < 0.001 generated by *t*-test.



Supplemental Figure S9. Tiller number and BR sensitivities of the representative overexpression plants. (A) Tiller number was counted at the harvesting stage. Bars indicate SD (n = 10). *P < 0.05 and ***P < 0.001 generated by t-test. (B and C) Coleoptile length and root length were measured after ten-day growth on agar media with or without BL. Bars indicate SEM (n = 10 - 19). *P < 0.01 and ***P < 0.001 generated by t-test.



Supplemental Figure S10. BR sensitivity of the overexpression plants evaluated by lamina bending test. HBL was applied at 3 days after germination when the 2^{nd} leaves were emerging and angles (the 2^{nd} leaves) were measured after additional 4-d-growth. Bars indicate SEM (n = 16 - 31). $^*P < 0.05$, $^{**}P < 0.01$ and $^{***}P < 0.001$ generated by t-test.

Supplemental Table S1. Primer sequences for mutation identification.

Name	5' - 3' sequence
GSK1-F	GCGATTTCTCATGCGATTTT
GSK1-R	ACCAGAACCTTTGCACTTCC
GSK2-F	GGTGCTCATTTTTGGTGGTT
GSK2-R	AGGGTGGAAAAGGCCAATAC
GSK3-F	GTGGGAACTGGATCATTTGG
GSK3-R	CCCTCTTGAAAACCAGAAAGAA
GSK4-F	GGACAAAGAATGTCGCATGA
GSK4-R	AGTAGCATCCGCCACAAATC
BZR1-F:	AGAGGGAAAGCACGCTACTG
BZR1-R:	CTCGGGAAGCTCGACGAC
BZR2-F:	GCGTGATCTCTCCGCTTC
BZR2-R:	AATCGCCCTAGCAAAGGATT
BZR3-F:	GACTGCTCCTCCTC
BZR3-R:	AGCTCGAATCGAAGCAAAAA
BZR4-F:	ATGATGAACGGAGGAGGGG
BZR4-R:	AGCGCCACAGCATCATCTCT
PPKL1-F	CTGCTGGTTTATCTGCCGAG
PPKL1-sg1-R	CAAACTTACATGCATGGTGGC
PPKL1-sg2-F	CAAACCACCGACGAATC
PPKL1-sg2-R	CCGTCCTCCTTCTCGAT
PPKL1-sg3-F	GTACAAAATTGTTTTCGGTT
PPKL1-sg3-R	GAGGCTCACCTTGTGGAGTAA
PPKL2-sg1-F	CCTGGTTGGTTTCAGC
PPKL2-sg1-R	GACGACAACGGCGAAGTAGA
PPKL2-sg2-F	GGAGCTAGGGTTTTGGAGGT
PPKL2-sg2-R	AAGGGATGAGGAGACCTGGA
PPKL3-sg1-F	TCCTAGGGTAACCCACCACT
PPKL3-sg1-R	TTGTTGGCACAAGCATGCAA
PPKL3-sg2/3-F	GGTGTGGCGTGTTCTACCTA
PPKL3-sg2/3-R	CCGTCCTCCTTCTCCAT
NPT-F	CATGTGTCACGACGAGATCC
NPT-R	CGCACAATCCCACTATCCTT
HPT-F	TAGGAGGCGTGGATATGTC
HPT-R	TACACAGCCATCGGTCCAGA

Supplemental Table S2. Primer sequences for vector constructions.

Name	5' - 3' sequence (Restriction enzyme sites were indicated)		
LC-PPKL2-F	GGCAGCGGCC <u>GAATTC</u> ATGGGGACGGCGGGAAG (EcoRI)		
LC-PPKL2-R	GTCGACTGCAGAATTCTCATATATAAGCAAGAGAAT (EcoRI)		
Flag-PPKL2-F	CGAAATCGAT <u>GGATCC</u> GATGGGGACGGCGGGAAG (BamHI)		
Flag-PPKL2-R	AGGCTACGTAGGATCCTCATATATAAGCAAGAGAAT (BamHI)		
LC-aGSK2-F	GGCAGCGGCC <u>GAATTC</u> ATGGACCAGCCGGCGC (EcoRI)		
LC-aGSK2-R	GTCGACTGCAGAATTCTTAGCTCCCAGTATTGAAGAAG (EcoRI)		
LM-GSK2-F	GGGCCGCGAC <u>TCTAGA</u> ATGGACCAGCCGGCGCC (XbaI)		
LM-GSK2-R	AAAGCAGGAC <u>TCTAGA</u> TTAGCTCCCAGTATTGAAGAAG (XbaI)		
Flag-GSK2-F	CGAAATCGATGGATCCGATGGACCAGCCGGCGCC (BamHI)		
Flag-GSK2-R	AGGCTACGTAGGATCCTTAGCTCCCAGTATTGAAGAAG (BamHI)		
LM-aOsBZR1-F	AAGAGGACTT <u>GAATTC</u> ATGACGTCCGGGGCGGCG (EcoRI)		
LM-aOsBZR1-R	CCGGGGTACC <u>GAATTC</u> TCATTTCGCGCCGACGCC (EcoRI)		
Flag-aOsBZR1-F	CGAAATCGATGGATCCTATGACGTCCGGGGCGG (BamHI)		
Flag-aOsBZR1-R	AGGCTACGTAGGATCCTCATTTCGCGCCGACGCC (BamHI)		
LC-OsBZR2-F	GGCAGCGGCC <u>GAATTC</u> ATGGCGACGGGAGGAGGAGG (EcoRI)		
LC-OsBZR2-R	GTCGACTGCAGAATTCTTAAGCAGCAGCTCTTGTCCTAGAGCT		
LC-OSDZR2-R	(EcoRI)		
For point mutation	For point mutation		
aOsBZR1-1F	ATGACGTCCGGGGCGGC		
aOsBZR1-1R	TCGCACTCCGGTATCGTGTCCAGGTGCTCGAGGCGGCGCC		
aOsBZR1-2F	GCCGCCGCCTCGAGCACCTGGACACGATACCGGAGTGCGA		
aOsBZR1-2R	TTTCGCGCCGACGCCGAGCG		

Supplemental Table S3. Primer sequences for RT-qPCR analysis.

Name	5' - 3' sequence
OsGSK1q-F	ACGGGTCACATCATCTCC
OsGSK1q-R	AGTTCCTACAACTCGCTCC
OsGSK2q-F	CCAACCCGTGAGGAAA
OsGSK2q-R	GCGTGAAGCGAGGTCTA
OsGSK3q-F	CAGCCTCTCTTTCCTGGTGA
OsGSK3q-R	GGGATTCATGCAGCGGATTT
OsGSK4q-F	GGAGGAAGAACGATG
OsGSK4q-R	TTGCCTGGAATACAACAC
D2q-F	AAACCGACATGGGTGAGACC
D2q-R	ACCGCAGCGTCTCTGTTATC
D11q-F	CCTAGGGGAGGACCACAAGA
D11q-R	CCAAGGTAGCTGGGCTTGAG
DWARF4q-F	GTGCTCAACTTCAGGTGGGA
DWARF4q-R	CCTAATGGGAAGGCCCTTGG
BRD2q-F	GCATTGCCGCCAAAAGTATG
BRD2q-R	TCCTGCACCTCCTTCTCAGTC
UBQq-F	GAGCCTCTGTTCGTCAAGTA
UBQq-R	ACTCGATGGTCCATTAAACC