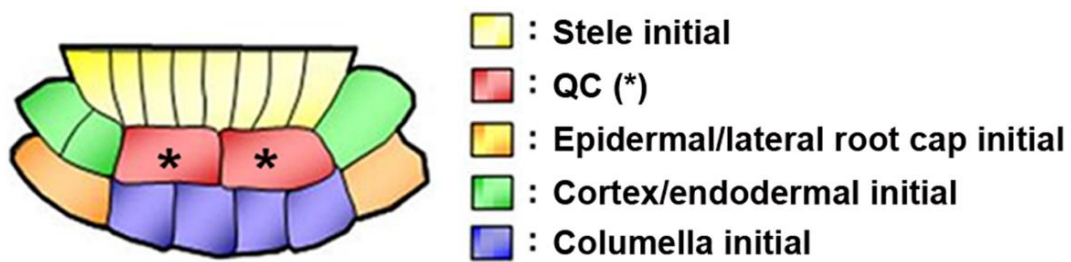
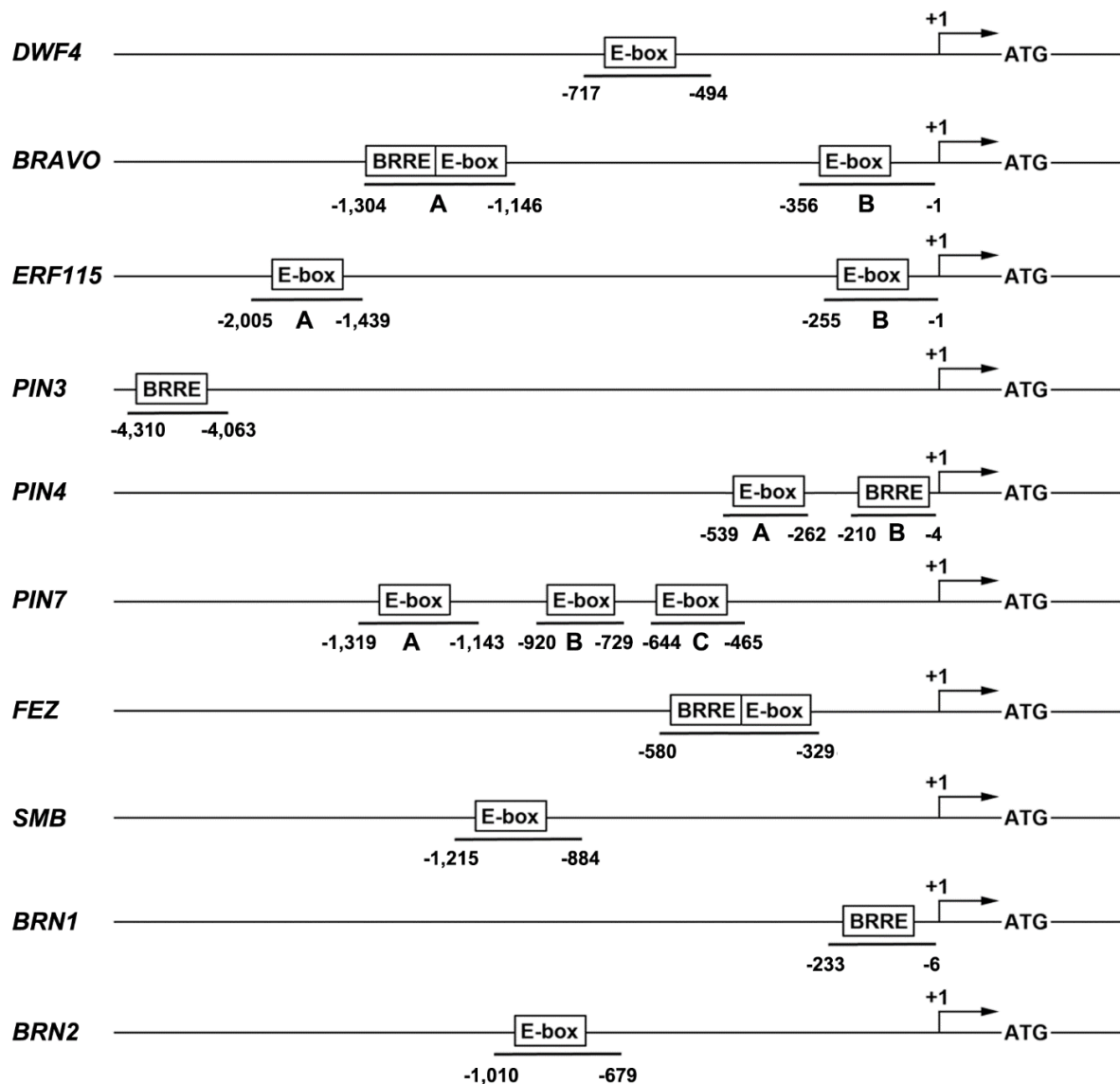


BZR1-dependent brassinosteroid signaling pathway leads to ectopic activation of quiescent cell division and suppresses columella stem cell differentiation

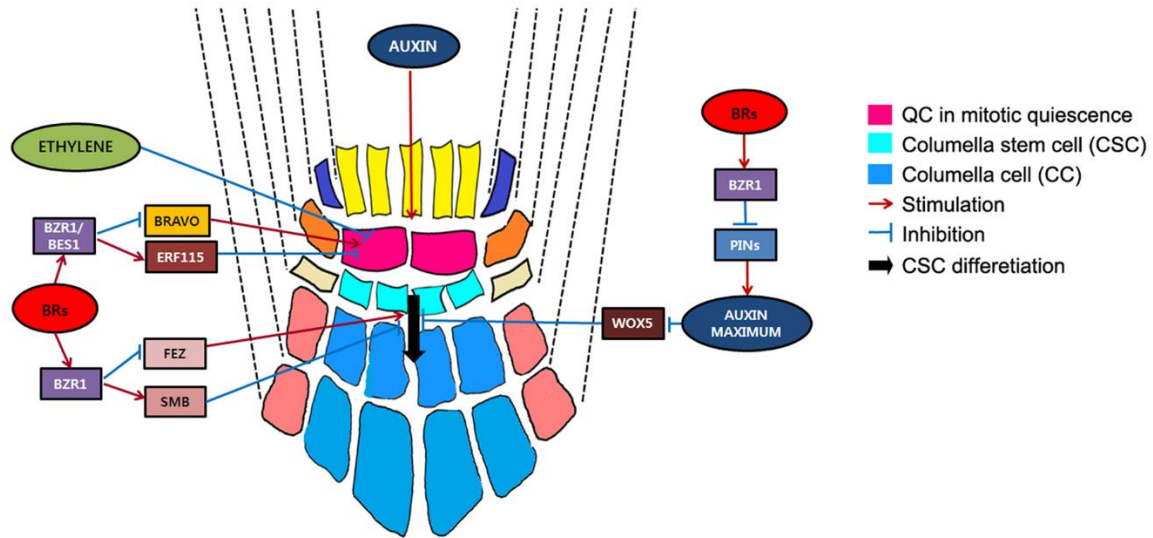
Hak-Soo Lee, Yoon Kim, Giang Pham, Ju Won Kim, Ji-Hye Song, Yew Lee, Yong-Sic Hwang, Stanley J. Roux, Soo-Hwan Kim



Supplementary Figure S1. Schematic diagram of the root SCN area.



Supplemental Figure S2. Schematic diagram of potential BZR1- and BES1- binding sites (BRRE and E-box) and the PCR-amplified DNA fragments used for ChIP-qPCR analysis. 5' and 3' position of each fragment is indicated by numbers. +1 represents the transcriptional initiation site of the corresponding gene. Nucleotide positions and fragments lengths are not in scale.



Supplemental Figure S3. A schematic model explaining brassinosteroid regulation of the QC maintenance and the columella stem cell (CSC) differentiation. Both BZR1-/BES1-mediated brassinosteroid signaling pathways promote mitotic reactivation of QC cells by inhibiting *BRAVO* (a negative regulator of QC division) and stimulating *ERF115* (a positive regulator of QC division) gene expression. In contrast, BRs regulate CSC differentiation into columella cells (CCs) in a BZR1-/BES1-dependent manner. BZR1-mediated inhibitory BR action on PINs expression and activities provoke proximal movement of auxin maximum leading to WOX5-mediated CSC differentiation into CCs. In other hand, BZR1 down-regulates gene expression of FEZ (encoding a protein promoting periclinal root cap-forming cell division), and this may result in another level of suppression for CSC differentiation into CCs. Ethylene pathway and BZR1-mediated BR signaling pathway acts independently on the QC reactivation.

Supplemental Table S1. Primers used in quantitative real time RT-PCR analysis		
Gene name	Locus	Primer set (5' to 3')
<i>WOX5</i>	At3g11260	TCCAACCTCCAAGGTGGACAAAATGA
		ATGGCGGTGGATGTTCCATTTCAG
<i>SCR</i>	At3g54220	TTCTCACCCCCTCACTGAGTTTTTG
		GTTGTTGGTCGTGAGATTGCATGG
<i>SHR</i>	At4g37650	TGGGAAGAGAGTTTTCCAAGGACGA
		TCATCCGCCACCTCATCACTATACC
<i>BRAVO</i>	At5g17800	CGAGGACACTGGAGACCAACAGAAG
		CAG CGT TAT CGG TAC GAC CTG GA
<i>ERF115</i>	At5g07310	CAAATCCGCAGACTAATCCGCAAAC
		AGGAGGTGAAGAATCCCCAAAACG
<i>PIN3</i>	At1g70940	GGAGCACCTGACAACGATCAAGG
		CTCGGCGTCTTTTGGTCTCTCTG
<i>PIN4</i>	At2g01420	ATGTGCATCCCACGATTCTAAGCAC
		CAATCTCCGAGGCTCTCTCAAAAGC
<i>PIN7</i>	At1g23080	TTGGGCTCTTGTTGCTTTCAGGT
		CCGCTGGTCCAGTAAAGAATCTCAC
<i>CUL3</i>	At1g26830	TCCCGATCATCCGTCTTTCCTCT
		GCAGGACCATGTTGTACGCATTTC
<i>ACS5</i>	At5g65800	GCGATGCTTTCCTTTTGCCTACTC
		TTTCTGGGCTTGTTGGTAAGCTTGT
<i>ETO1</i>	At3g51770	TGCTGGATGCAGCTGTATGATCGTT
		GCTGCCTTTTGACAATTGAGCCGTA
<i>FEZ</i>	At1g26870	TCAGTTTGCAGCACCTTCATGTTTC
		CACCGTGGATGACGACCCTATCT
<i>SMB</i>	At1g26870	TCGGAAATGGGAGATAGAAACAACG
		TTGGAAGATCCCAGGGGTCATATTT
<i>BRN1</i>	At1g33280	CACGTGCAAGGCAGTAAGTGAATGG
		TCACCGCGCAATGAAAGCAGATTAG
<i>BRN2</i>	At4g10350	TCAAGCCAACCCTAGTGAAGATGGA
		TGGACTGTCTCGGTGCATAAAGCTA
<i>CPD</i>	At5g05690	GCGGTGTTTTTCAGACGTGCAAT
		GAAAGTGCGAGCATCTTTGAAGTGG

Supplemental Table S2. Primers used in ChIP-qPCR analysis		
Gene name	Locus	Primer set (5' to 3')
<i>UBC30</i>	At5g56150	CAAATCCAAAACCCTAGAAACCGA
		AACGACGAAGATCAAGAACTGGGAA
<i>PP2A</i>	At1g69960	AGCAGCACAACCCTCAACAG
		CCAGATGTGCTAAAGACGGAG
<i>DWF4</i>	At3g50660	GTGTTTTCTGACTATTGAGGGG
		CGGTACGGTCTCAATCGGTTTA
<i>BRAVO A</i>	At5g17800	AAAATTTAAATTTAAACTAGTAGCAAAAAAT
		TTTTACTTATATACTATATTCAGTG
<i>BRAVO B</i>	At5g17800	AAAAAAAAAAAAAAAAATGATAAATAAAA
		GAGAGCACTTGAATGGCTTTTCACTG
<i>ERF115 A</i>	At5g07310	CGTTCTCGTCAACAAATCTGAAAATAC
		CAATCGAGAACTGTTGTCTTTTTTTT
<i>ERF115 B</i>	At5g07310	TATGCAAACTTCTGCTTGACGTAA
		CTTTGCTAAAATCTTTAAACCTCTTT
<i>PIN3</i>	At1g70940	CTCCAATACTCGATCGTGAAGA
		GGATGATAGAGTGTGGATTGG
<i>PIN4 A</i>	At2g01420	CAAAAACAAAAACAAAATAT
		AAAGTTGCAAAGGAACCTTG
<i>PIN4 B</i>	At2g01420	GCACGACTATTCCATAAACTGT
		GGATTCGGTGAAGAGGACTA
<i>PIN7 A</i>	At1g23080	CCAAACCATGAGCAGAATTGT
		CGTTTACACAATTATAATAGCAG
<i>PIN7 B</i>	At1g23080	CGGTCGCGGAAAGATCTTGA
		CCCAAGAAATCTCACTTTTAAG
<i>PIN7 C</i>	At1g23080	CGGCGAATATGATCTTGCATT
		GGGCCACTTAACGTATTACTAG
<i>FEZ</i>	At1g26870	CGATCGGCCCTTGTATCCTTTTAT
		AAAAATGCACTATCATCATATCGT
<i>SMB</i>	At1g26870	TAAGGAGCAAATATGGAACCTTA
		AGGATTAACATACTAAGCAGTAA
<i>BRN1</i>	At1g33280	CTCCGGCGGGCGTTGTCACCGGCCG
		GTTAATAAATGATCAATGTTTGTTTG
<i>BRN2</i>	At4g10350	TACATTAATATAACACATATTATT
		AGAAACACAAAAAGAAAAACCTTT

Supplemental Table S3. Division of QC in various brassinosteroid mutants

	Col	<i>bril-116</i>	<i>bzrl-1D</i>	<i>bril-116</i> <i>/bzrl-1D</i>	En-2	<i>bes1-D</i>
DAG 3	0/25 (0) ^a	0/32 (0)	0/24 (0)	0/28 (0)	0/25 (0)	0/31 (0)
DAG 7	2/38 (5)	0/26 (0)	14/28 (50)	10/27 (37)	0/23 (0)	21/25 (84)
DAG 14	5/28 (18)	2/22 (9)	24/32 (75)	20/29 (69)	2/21 (10)	33/33 (100)

^a: Number of plants with periclinal division of QC / number of plants observed (percentage of plants with the reactivated QC). DAG: Day after germination.