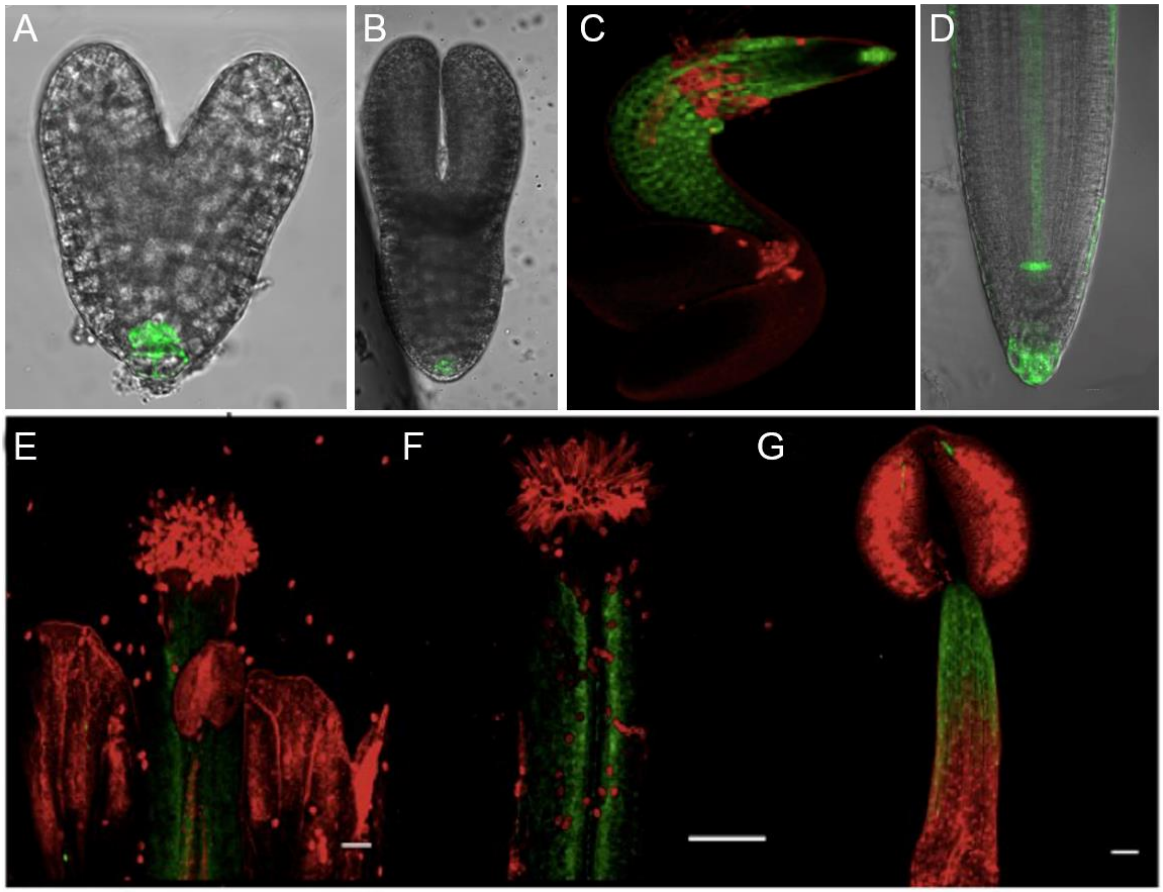

Figure S1. *At3g08030* expression during *Arabidopsis thaliana* development using *pAt3g08030::ER-GFP* plants. A) GFP fluorescence in embryo at heart stage. B) GFP fluorescence in embryo at torpedo state. C) GFP fluorescence in hypocotyl and primary root of a 36-h germinating seed. D) GFP fluorescence in the primary root from 7-d seedlings. E) GFP fluorescence in flower. F) GFP fluorescence in carpel and G) GFP fluorescence in stamen. *pAt3g08030::ER-GFP* transgenic plants were obtained and laser confocal scanning microscopy was done as described in Salazar-Iribe *et al.* [19]. ER-GFP refers to Endoplasmic Reticulum signal peptide fused to GFP, A, B, D are longitudinal sections and C, E, F and G are projections of confocal stacks.

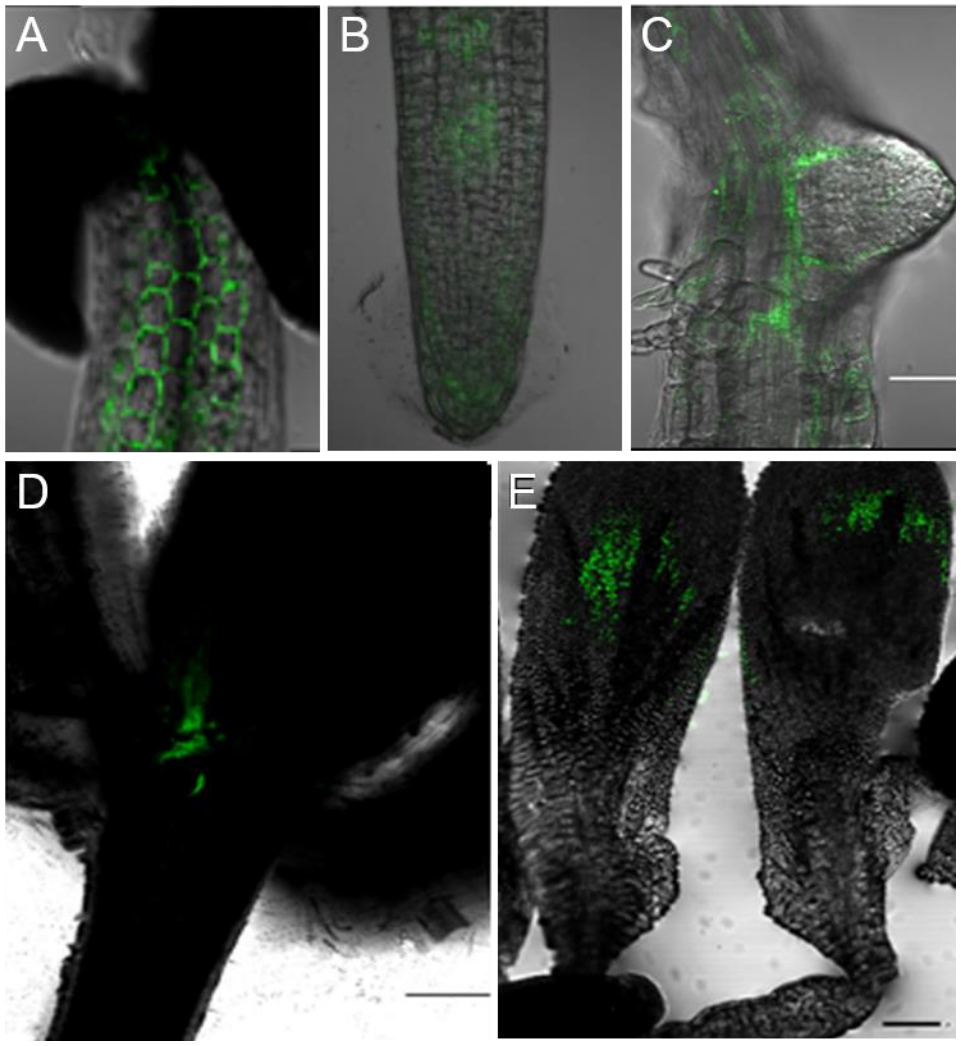

Figure S2. *At5g11420* expression during *Arabidopsis thaliana* development using *pAt5g11420::ER-GFP* plants. A) GFP fluorescence in hypocotyl of a 48-h germinating seed. B) GFP fluorescence in the primary root from 7-d seedlings. C) GFP fluorescence during lateral root emergence. D) GFP fluorescence in flower. E) GFP fluorescence in petals *pAt5g11420::ER-GFP* transgenic plants were obtained and laser confocal scanning microscopy was done as described in Zúñiga-Sánchez *et al.* [13]. Images are projections of confocal stacks.

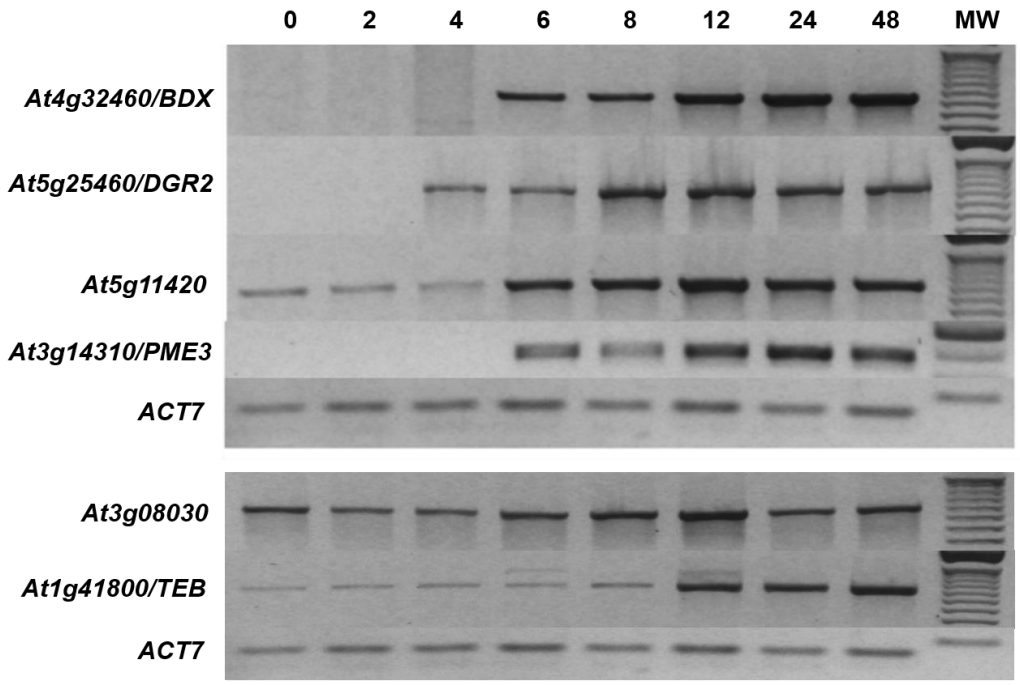

Figure S3. *At4g32460*/*BDX, At5g25460/DGR2* , *At5g11420*, *At3g14310/PME3* , *At2g41800/TEB* and *At3g08030* expression during seed germination. Seeds were collected after 2, 4, 6, 8, 12, and 24 h of imbibitions. 0 corresponds to dry seed and 48 to germinated seeds. *ACT7* expression was used as control. Primers used: BDXF 5´GTGATAGTGCTTCTTCTCCTTCAC 3´; BDXR 5´AGCGACGAATCTCAATGAC 3´; DGR2F 5´CTTCCTTCTTTTCATCGCC 3´; DGR2R 5´ACGAGAAATCATCGCTCC 3´; At5g11420F 5´AATCGCCACCATCACTTC3´; At5g11420R 5´ CATAACACTTGTGCGGGTC; 3´At3g08030F 5´GGTTCCCAAAGCCATTATTC 3´; At3g08030R 5´ ACAATCTCGTCAATGACAGG 3´; ´TEBF 5´TCCTCCTCCTATCTCTCTGC 3´; TEBR 5´AAACGGTTCTCTTCCTGC 3´; PMEF 5´CATCAATGAAAGAAATTTTTTC 3; PMER 5´AGACCGAGCGAGAAGGGGAAA 3´; ACT7F 5´GGTCGTACAACCGGTATTGT 3´; ACT7R 5´GAAGAGCATACCCCTCGTA 3´. PCR analyses were done as described in Garza-Caligaris *et al.* [36].

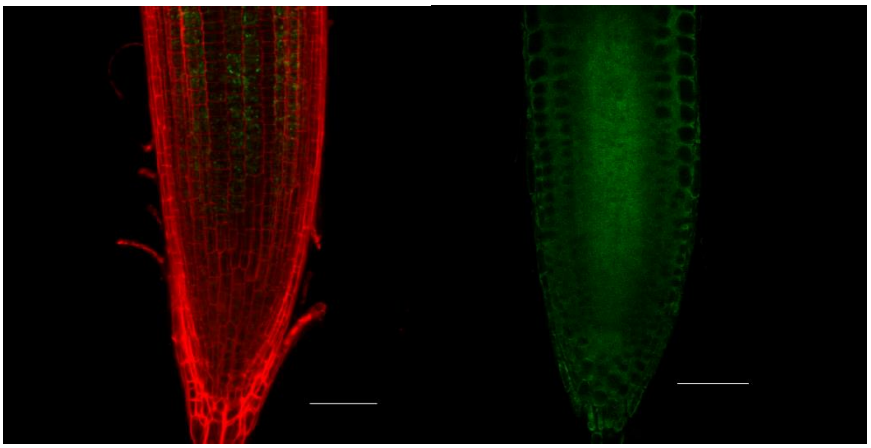

Figure S4. Subcellular localization of BDX in *Arabidopsis thaliana* primary roots under salinity stress conditions using *pBDX::BDX-GFP* plants. A) GFP fluorescence is detected intracellulary in the epidermal cells of the primary root from seedlings grown under control conditions. B) GFP fluorescence is detected in the cell wall of the epidermal cells o f the primary toot of seedlings grown under salinity stress.

Seedling were grown in MS with 50 mM NaCl. *pBDX::BDX-GFP* plants transgenic plants were obtained and laser confocal scanning microscopy was done as described in Salazar-Iribe *et al.* [30]. A is a longitudinal section and B is a projection of confocal stacks.

References for Supplemental material

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