

**Investigating the molecular mechanism
whereby auxin modulates *Arabidopsis thaliana*
growth under salinity stress conditions**

by

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An RT-qPCR experiment was conducted using *EXPA11* qPCR primers on cDNA synthesised from RNA extracted from No-0 and 35S::*EXPA11* T3 plants grown on soil under standard conditions. Four leaves were pooled per plant for RNA extraction for each of the genotypes. The 35S::*EXPA11* T3 lines that were screened are ‘2,2-4’, ‘7,1-3’, ‘9,2-5’, ‘18,1-2’, ‘20,1-3’, and ‘21,1-3’. *EXPA11* expression was normalised to the reference gene *MON1*.

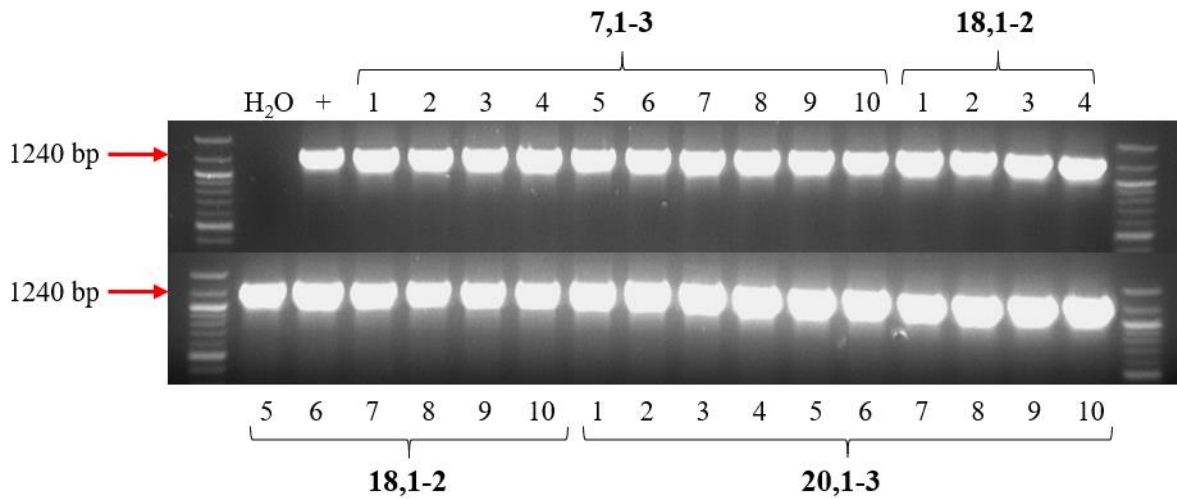


Figure 7.6: Presence of 35S::*EXPA11* construct in T3 lines

For each third transgenic generation (T3) line, DNA extractions were performed on 10 seedlings. The PCR with p35S F and *EXPA11-attB2* R primers was conducted to confirm the presence of the 35S::*EXPA11* construct in the genomic DNA (gDNA) of the T3 lines. The T3 lines that were screened are ‘7,1-3’, ‘18,1-2’, and ‘20,1-3’. The positive control used the gDNA from the second transgenic generation (T2) of the T3 line 18,1-2 as template. The MW marker included is the New England Biolabs (NEB) Quick-Load® 100 bp DNA ladder (Catalogue #NO467S).

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