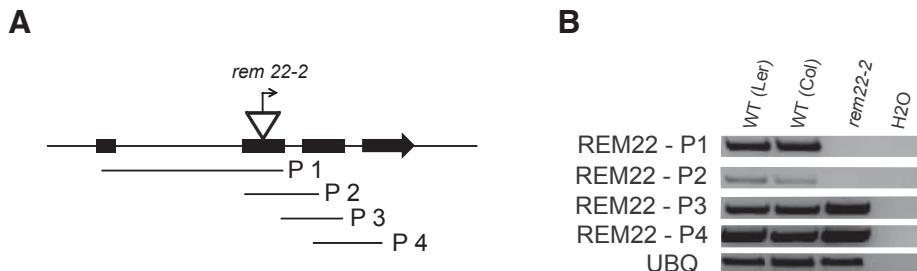


SUPPLEMENTARY MATERIAL

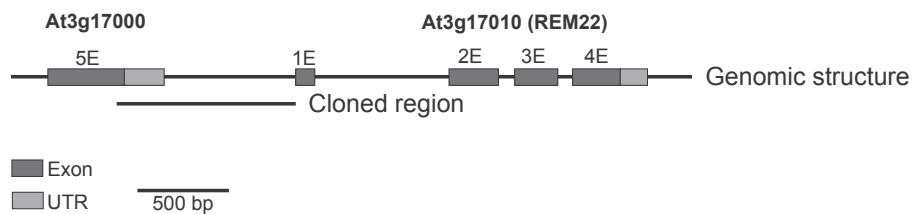
corresponding to:

Reproductive Meristem22 is a unique marker for the early stages of stamen development

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ELLIOT MEYEROWITZ and MARCIO ALVES-FERREIRA



Supplementary Fig. 1. Position of the T-DNA insertion in the REM22 gene and gene expression analysis. **(A)** The triangle indicates the location of T-DNA insertion in the second exon of REM22 gene. The lines below the gene diagram show the PCR fragments produced by distinct primer combination used in the RT-PCR analysis (Supplementary Table 1). Exons are represented as black solid boxes. **(B)** REM22 expression in inflorescence of distinct ecotypes (Ler - Landsberg erecta; Col - Columbia) and in the insertion line (SALK_091149). Gene-specific primer combinations for REM22 used in the RT-PCR are indicated in the left side. UBIQUITIN10 (UBQ) was used as a control for loading. Complete REM22 transcripts were only detected in WT plants.



Supplementary Fig. 2. Genomic structure of REM22 gene. REM22 gene, the promoter region and the 5th exon of the gene At1g17000 are shown. The cloned promoter region (1022 bp upstream the REM22 ATG) is indicated by the black line below the gene diagram. The exons (dark gray), the UTRs (light gray) and the scale bar in base pairs (bp) are shown below the diagram.

SUPPLEMENTARY TABLE 1

PRIMER LIST

Gene – RT-PCR fragment	Forward	Reverse
REM22 – P1	GGCAGATCTATCCTCTGACTG	CGAGGTTACGCATCTCTAAGC
REM22 – P2	GGAAAGACATGGGATGAGTC	TCTATGAGGCTGTTCTCTCCCT
REM22 – P3	GGGTCTTCTAGGGATCAAG	GGTATTGCCAGGAACTTGAGGT
REM22 – P4	GGACGAGCTGAGTCCTAGTTAGA	AGATTGTCGAAATCTCCCTCAC
UBQ10 (AT4G05320)	GATCTTGCAGAAACATTGGAGG	CGACTTGTCAATTAGAAAGAAAGAGAT