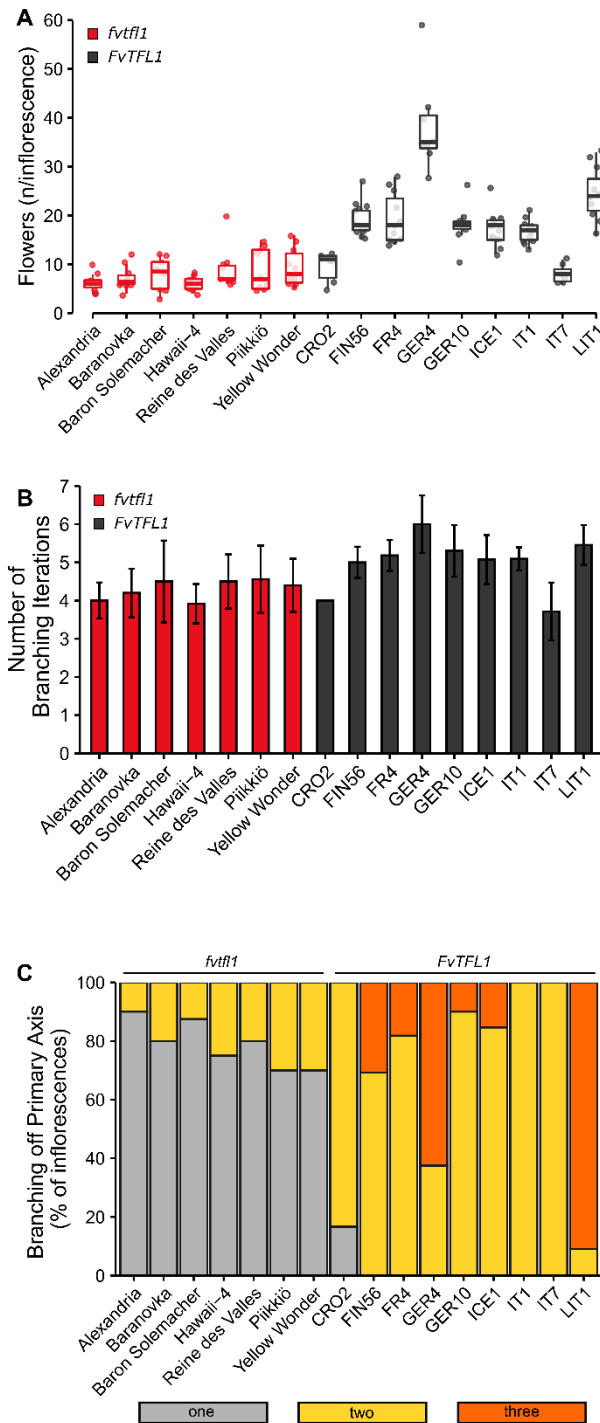
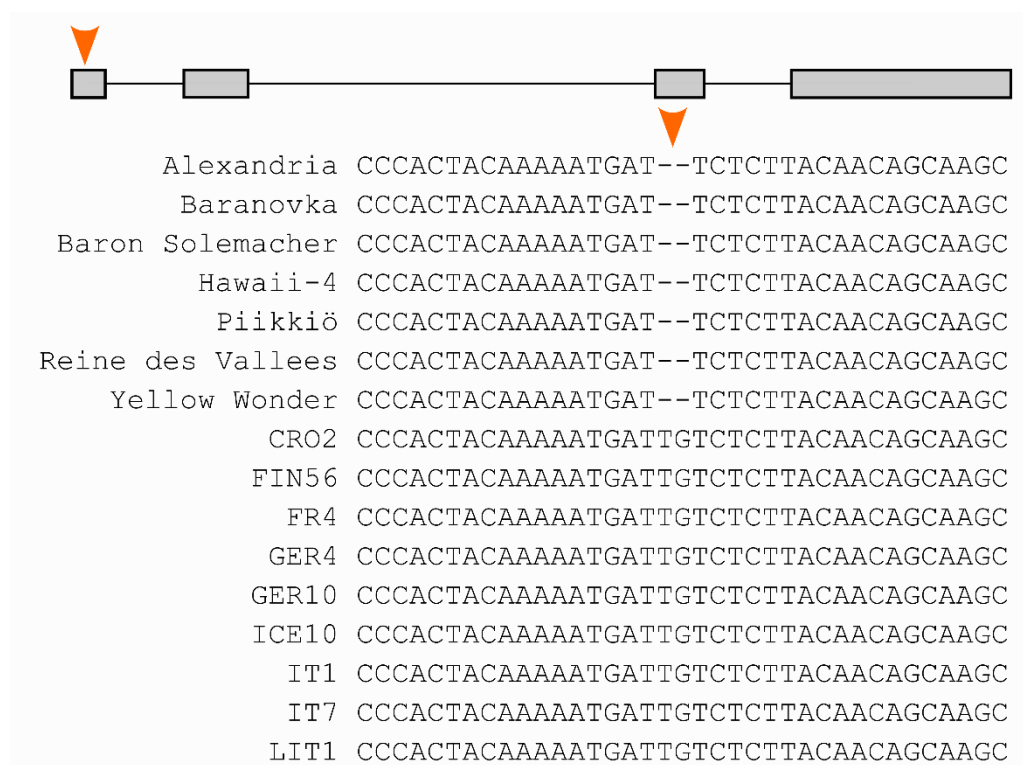


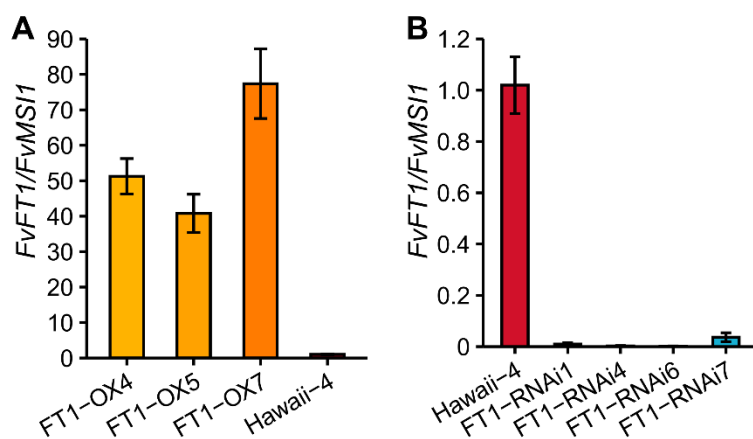
Figures



**Supplemental Figure S1.** Inflorescence phenotypes of seven *fvtfl1* and nine *FvTFL1* genotypes. Supports Figure 3. **A.** Number of flowers on the first emerged fully-developed inflorescence per plant (n = 6 – 13). Boxplots and points show distribution of raw data. Each point represents an individual inflorescence. **B.** Average number of branching iterations along the longest branching path. Bars and error bars represent the mean ± standard deviation (n = 6 – 13). **C.** Percentage of observed inflorescences with one, two, or three branches on the primary branching axis.



**Supplemental Figure S2.** Alignment of the part of the first exon of *FvTFL1* gene in seven cultivars and nine WT accessions. Supports Figure 3.



**Supplemental Figure S3.** *FvFT1* expression levels in the leaves of *Hawaii-4* (control), *FvFT1* overexpression (OX) and *FvFT1* silencing (RNAi) transgenic lines. Supports Figure 4. Error bars and show standard error ( $n = 3 - 4$ ). Biological replicates represent 1cm  $\varnothing$  leaf disc excised from a plant at ZT16. Expression levels were normalized to *Hawaii-4*. *FvMSI1* was used as a reference gene.

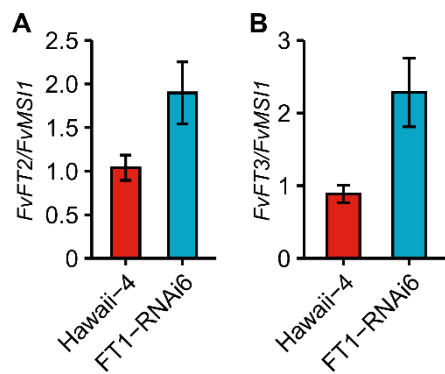


**Supplemental Figure S4.** *FvFT1* overexpression in the *fvtf1* (*Hawaii-4*) background induces flowering in all axillary buds and reduces overall plant size. Supports Figure 4. Scale bar equals 5 cm.



**Supplemental Figure S5.** *FvFT1* overexpression in the functional *FvTFL1* background. Supports Figure 4. **A.** Overall plant phenotypes. **B.** The number of flowers per inflorescence in *FvFT1* overexpression plants in functional and non-functional *FvTFL1* backgrounds. Boxplots and points show the distribution of raw data. Each point represents an individual inflorescence ( $n = 16 - 54$ ). Up to six inflorescences per plant were examined. Data were analyzed by fitting a generalized linear model (GLM). \*\*\* indicate  $P$  value  $< 0.0001$  between the groups (Tukey's HSD).

Supplemental Data. Lembinen et al. (2023). Plant Cell.



**Supplemental Figure S6.** The expression of *FvFT2* (A) and *FvFT3* (B) in the FMs of *Hawaii-4* and *FT1-RNAi6* plants. Supports Figure 4. Bars and error bars show mean  $\pm$  standard error ( $n = 3 - 4$ ). Expression levels were normalized to *Hawaii-4*.

## Tables

**Supplemental Table S1.** Model parameters used to produce inflorescences in Figure 7 (Phenotype 0) and Figure 8 (Phenotypes 1-7). Parameters shown in bold are discussed in the main text or Supplemental Model Description; the remaining parameters are explained in the code. Only values differing from the previous column are shown. The key parameter (in red) differentiating the models is `veg_half_life`, which determines the time (measured in plastochrons) over which `veg` decreases to  $\frac{1}{2}$  of its initial value. Related parameters are grouped together as in the code.

Phenotype	0	1	2	3	4	5	6	7
max_age	8							
dt	0.02							
plastochron	1							
delay_mono	0.44							
delay_sym	0.3	0						
delay_diff								
veg_init	1							
<b>veg_half_life</b>	<b>1.98</b>	<b>2.77</b>	<b>2.42</b>	<b>2.07</b>	<b>1.64</b>	<b>1.37</b>	<b>1.05</b>	<b>1.05</b>
th_m	0.43							
th_s	0.27				0.23			0.6
th_diff	0.08				0			
b_th_1	0.6						0.48	
b_th_2	0.45							
pleiotropic_scale	4	1				0.82	0.5	0.4
peduncle_length_ctrl	0.4	0.2		0.3	0.65	0.7	0.75	0.6
int_len_power	1.2							
internode_segments	10							
distal_diameter	0.011				0.008	0.011		
pipe_model_power	3							
angle_main	12	35	38	17	27			35
angle_lateral	10	17	13	29	7			17
angle_sympodial	20	32			35		32	
int_elasticity	0.32	0.06						
int_elasticity_power	1.3							
bract_angle_branch	5							
bract_angle_no_branch	25							17
ang_adjustment_1	0						6	14
ang_adjustment_2	35							
petiole_length_1	0.2	0.7					0.3	0.3
petiole_length_2	0.02							
petiole_elasticity_1	1.8							
petiole_elasticity_2	1							
bract_size_power	0.5							

**Supplemental Table S2.** Plant material used in this study.

<b>Accession Name</b>	<b>Origin/Collection Site</b>
<i>Alexandria (PI602923)</i>	NCGR*
<i>Baranovka</i>	Lyubov Kuznetsova**
<i>Baron Solemacher (PI551507)</i>	NCGR*
<i>Hawaii-4 (PI551572)</i>	NCGR*
<i>Piikkiö</i>	Unknown
<i>Reine des Vallees (PI551824)</i>	Dr. Amparo Monfort***
<i>Yellow Wonder (PI551827)</i>	NCGR*
<i>CRO2</i>	44°46'15.1896" (N) 15°38'51.6408" (E)
<i>FIN56 (PI551792)</i>	NCGR*
<i>FR4</i>	48°25'24.4452" (N) 7°39'48.0168" (E)
<i>GER4</i>	50°59'6.9036" (N) 11°19'21.216"(E)
<i>GER10</i>	47°33'21.168" (N) 10°1'17.2632" (E)
<i>ICE10</i>	64°45'36.774" (N) -21°35'35.2824" (E)
<i>IT1</i>	45°56'13.362" (N) 10°48'20.2392" (E)
<i>IT7</i>	46°16'30.1188" (N) 11°16'54.9372" (E)
<i>LIT1</i>	54°47'51.7812" (N) 25°20'54.2292" (E)

\*National Clonal Germplasm Repository, Corvallis, USA; \*\* The Federal Research Center Institute of Cytology and Genetics, Novosibirsk, Russia; \*\*\* Centre for Research in Agricultural Genomics, Barcelona, Spain

**Supplemental Table S3.** Primers used in this study.

<b>Name</b>	<b>Forward</b>	<b>Reverse</b>	<b>Reference</b>
<i>FvTFL1 (FvH4_6g18480)</i> (cloning/sequencing)	ATGGCAAGAATGTCGGA ACC	CTAGCGTCTTCTTGCT GCC	
<i>FvTFL1 (FvH4_6g18480)</i> (RT-qPCR)	AACGGCAGCAACAGGAA C	CTGGCACCACAGATGC TACA	Koskela et al., 2012
<i>FvAP1 (FvH4_4g29600)</i> (RT-qPCR)	AGCTCAGGAGGTTTCATG ACTG	TAAGGTCGAGCTGGTT CCTC	Koskela et al., 2012
<i>FvMSI1 (FvH4_7g08380)</i> (RT-qPCR)	TCTCCACACCTTTGATT GCCA	ACACCATCAGTCTCCT GCCAAG	Mouhu et al., 2009
<i>FvFT1 (FvH4_6g00090)</i> (RT-qPCR)	CAATCTCTTGGCCGAAA ACT	TGAGCTCAAACCTTCC CAAG	Koskela et al., 2012
<i>FvLFYa (FvH4_5g09660)</i> (RT-qPCR)	GATGACAGAATCAATGG AGGAG	CTGGTTTGTGACCTTG GTGG	
<i>FvFT2 (FvH4_4g30710)</i> (RT-qPCR)	ACTCGGTGGCTTGTGTT TTC	ATCACTCTCCCGACGA CAAG	Nakano et al., 2015
<i>FvFT3 (FvH4_3g09870)</i> (RT-qPCR)	AGCCGTTACCAAGTCT GTG	GTGGACAACATGAGAA GGTTTG	Nakano et al., 2015