

1 **Supplementary Information**

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3 **The floral repressor *GmFLC-like* is involved in regulating flowering time**
4 **mediated by low temperature in soybean**

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6 **This PDF file includes:**

7 Supplementary Fig. 1-2

8 Supplementary Table 1

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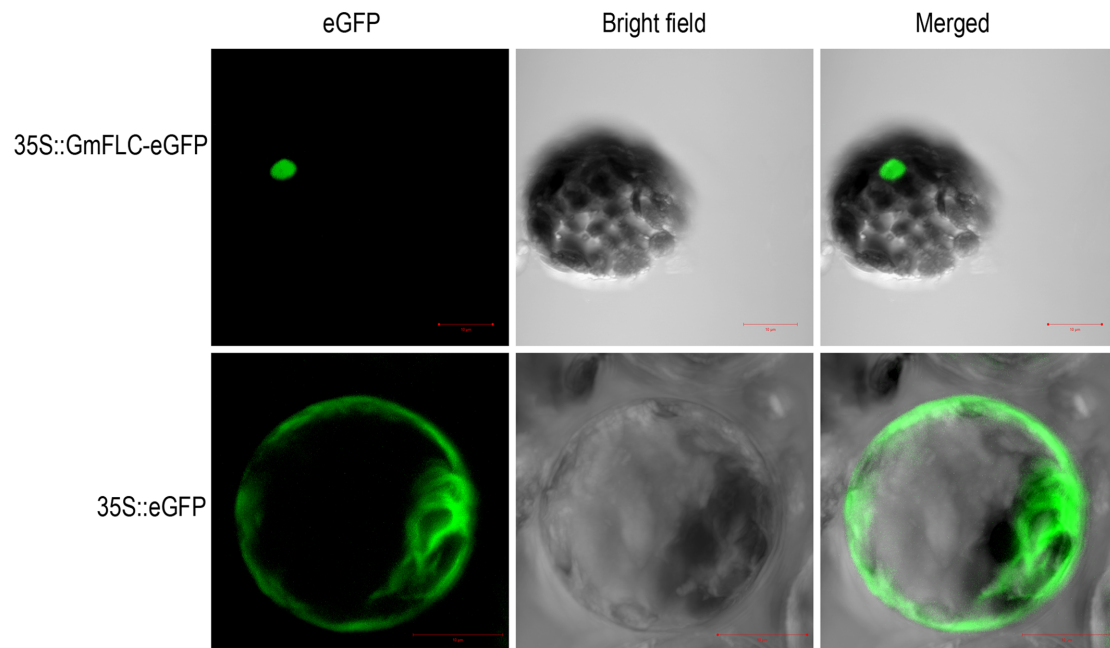
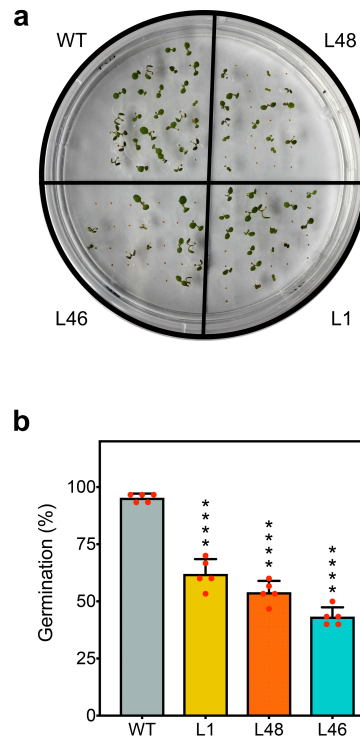


Fig. S1: Subcellular localization of GmFLC-like protein in Arabidopsis protoplast cells.

Constructs 35S::*GmFLC-like-eGFP* were transformed into Arabidopsis protoplast cells. The Empty plasmid 35S::*eGFP* was used as a control. The cells were observed under a confocal laser microscope. Scale bars, 10 μ m.



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24 **Fig. S2: Comparative analysis of seed germination rate among four lines of Arabidopsis. a**

25 Germinating status of WT (Col-0) and three *GmFLC-like* transgenic lines L1, L46, and L48 seeds

26 on 1/2 MS medium at 3 days after transfer to light. **b** Seed germination rate of WT and three

27 transgenic lines on 1/2 MS medium. Germination percentage was counted for approximately 30

28 seeds for each line. Experiments were repeated five times and mean value $\pm$ SD is plotted on the

29 graph. Significant differences according to the *t*-test are denoted as follows: * $p < 0.05$, ** $p <$

30 0.01, *** $p < 0.001$, **** $p < 0.0001$.

Supplementary Table 1. Primers used in this study

Primers	Sequence
For gene isolation	
<i>K-FLC</i> -F	ATGGGGAAGAAGAAGCTGGAGAT
<i>K-FLC</i> -R	CAACCTCTACCACTAGGCCAATCAT
For vector construction of overexpression in Arabidopsis	
<i>1301-FLC</i> -F	CGCGGATCC(BamHI)ATGGGGAAGAAGAAGCTGG
<i>1301-FLC</i> -R	CGGGGTACC(KpnI)TTATTTATTTATACTGAGTTCAAGAATT
For vector construction of subcellular localization	
<i>FLC-GFP</i> -F	CGGGGTACC(KpnI)ATGGGGAAGAAGAAGCTGG
<i>FLC-GFP</i> -R	CGGGATCC(BamHI)TTTATTTATACTGAGTTCAAGAATTGAG
For gene promoter isolation	
<i>Pro-GmFLC</i> -F	TTGCTTCGGTTACTGTTTCTTCC
<i>Pro-GmFLC</i> -R	TGTTCTCGATTCGCTTTATCTCC
<i>proFT2a-1-F</i>	ATTGGTACCCCGGGTGGGGAAGGGCTACT
<i>proFT2a-1-R</i>	ATTCTCGAGAACATTTCCCTCCCTTCTC
<i>proFT2a-2-F</i>	ATTGGTACCTCCTTTTTTCACTCAAGTG
<i>proFT2a-2-R</i>	ATTCTCGAGTTACTTATTTAATGGAAACTA
<i>intFT2a-1-F</i>	ATTGGTACCTATGATTTTAGTTTCATT
<i>intFT2a-1-R</i>	ATTCTCGAGTAATGGATGCTATATCAT
<i>intFT2a-2-F</i>	ATTGGTACCTTATTTATCTATCTCTTTT
<i>intFT2a-2-R</i>	ATTCTCGAGTGACTTTAAGTCCTATAAAA
For transgenic plants confirmation	
<i>Hpt</i> -F	ACTTCTACACAGCCATCGGTCC
<i>Hpt</i> -R	AGCGAGAGCCTGACCTATTGC
For qRT-PCR analysis	
<i>Gm-FLC</i> -F	TGACGCATAATCTGCTCCCTG
<i>Gm-FLC</i> -R	GCTAAACCATGGCATAGTTCCCT
<i>Gm-β-Tublin</i> -F	CCTCGTTCGAATTTCGCTTTTTTG
<i>Gm-β-Tublin</i> -R	CAACTGTCTTGTCGCTTGGCAT
<i>GmFT1a</i> -F	CCTTTTACACCCTGGTTATGG
<i>GmFT1a</i> -R	CCTGGAGGTTGCAGAGTTAGT
<i>GmFT1b</i> -F	GACTTCAGGACCTTTTACACCC
<i>GmFT1b</i> -R	GCTCACAACCTCTTCACCGA
<i>GmFT2a</i> -F	ATCCCGATGCACCTAGCCCA
<i>GmFT2a</i> -R	ACACCAAACGATGAATCCCCA
<i>GmFT2b</i> -F	GACATTCCAGCAACAACGG
<i>GmFT2b</i> -R	ATAGCCTTCTTCCACCACAAC
<i>GmFT3a</i> -F	GGATTCATCGTTTCGTGTTTG
<i>GmFT3a</i> -R	CACCAGAGCCAGTTTCCCT
<i>GmFT3b</i> -F	CTATGAAAGCCCACGACCC
<i>GmFT3b</i> -R	TTGAAGAAGACAGCAGCAACC
<i>GmFT4</i> -F	GTGAGTTCAAACCTTCCCAAAT
<i>GmFT4</i> -R	CAATCCGATGAATCCAGAA
<i>GmFT5a</i> -F	ACAGATTATGGTAGCAACGGAA
<i>GmFT5a</i> -R	CAAGGATAGCCAGAAAAGAAAG

<i>GmFT5b</i> -F	CTCAATCCTTTTACAATCTCCG
<i>GmFT5b</i> -R	CCTTAGGTCTTCACCACCAACA
<i>At-FT</i> -F	CCCTGCTACAACCTGGAACAAC
<i>At-FT</i> -R	AAGAACAAGGTAACCCAATGAAC
<i>At-SOC1</i> -F	AAACGAGAAGCTCTCTGAAAAG
<i>At-SOC1</i> -R	AAGAACAAGGTAACCCAATGAAC
<i>At-API</i> -F	GCAAGCAATGAGCCCTAAAG
<i>At-API</i> -R	ACTGCTCCTGTTGAGCCCTA
<i>At-TUB2</i> -F	ATCGATTCCGTTCTCGATGT
<i>At-TUB2</i> -R	ATCCAGTTCCTCCTCCAAC
