

Fig. S1. FLC is down regulated in the top $1 \alpha$ mutant. The qRT-PCRs were used to measure the relative expression levels of $F P A, F Y, F C A, F V E, F L K, F L D, L D, V R N 2$, VIP4, FLC, FRI, FT and SOC1 in topl $\alpha$ mutants. The expression levels of these genes in the wild-type Col- 0 were normalized to 1 and marked with the black line. Mean $\pm$ SD with three independent biological replicates. Bars marked with different letters are statistically different to each other ( $\mathrm{P}<0.05$ by Student's t test). Total RNA was isolated from 8-day-old seedlings grown in $1 / 2 \mathrm{MS}$ media under long-day conditions. TUBULIN was used to normalize the mRNA levels. The primer sequences used are listed in Table S1.




Fig. S2. The pTOP1a::TOP1a-GFP transgenic plants rescue early flowering phenotypes of top $1 \alpha$ mutants. (A) The early flowering phenotypes of top $1 \alpha$ mutants were rescued by the $p T O P 1 \alpha:: T O P 1 \alpha-G F P$ transgene. Bar, 2 cm . (B-C) The rosette leaf numbers and days of flowering were measured in Col-0, topla and pTOP1 $::$ TOP1 $\alpha$-GFP/top1 $\alpha$ transgenic plants. Mean $\pm$ SD. Bars marked with different letters are statistically different to each other ( $\mathrm{P}<0.05$ by Student's t test). Plants were grown in the soil under long-day conditions. $n$, the number of plants.


Fig. S3. TOP1 $\alpha$ expression patterns in the shoot apex. (A-E) TOP1 $\alpha$ expression patterns in the Col-0 seedlings using in situ hybridization. Serial sections were shown. Bar, $100 \mu \mathrm{~m}$. ( $\mathbf{F}$ ) The sense control of TOP $1 \alpha$ in the in situ hybridization. (G-I) The protein localization of TOP $1 \alpha$ in the $p T O P 1 \alpha::$ TOP1 $\alpha-G F P /$ top $1 \alpha$ rescued plant. Bar, $100 \mu \mathrm{~m}$.


Fig. S4. TOP1a does not regulate SDG8 and CTD at the transcription level. The relative expression levels of $S D G 8(\mathbf{A})$ and $C T D(\mathbf{B})$ in top $1 \alpha$ mutants were detected by qRT-PCR. Mean $\pm$ SD with three independent biological replicates. Bars marked with same letters are not statistically different to each other ( $\mathrm{P}>0.05$ by Student's t test). Total RNA was isolated from 8 -day-old seedlings grown in $1 / 2$ MS media under long-day conditions. TUBULIN was used to normalize the mRNA levels. The primer sequences used are listed in Table S1.


Fig. S5. TOP1 $\alpha$, CTD and SDG8 are in the same complex. BiFC-based FRET was performed to examine the interaction of TOP1 $1 \alpha$, CTD and SDG8 proteins in the same complex. The $35 S::$ CFP-TOP1 $\alpha-1$, $35 S::$ nYFP-CTD and $35 S::$ cYFP-SDG8 were co-transformed into the abaxial leaves of N. benthamiana. The 35S::cYFP-AGO6 and $35 S:$ :WOX5-YFP were used as negative controls. The excitation wavelength of 448 nm (CFP) and the emission wavelength of 514 nm (YFP) were used in the FRET channel. Bar, $100 \mu \mathrm{~m}$.


Fig. S6. The alignment of Topoisomerase 1 in different species. (A) All protein sequences were obtained from NCBI (https://www.ncbi.nlm.nih.gov/) and aligned using the software ClustalX2 (http://www.clustal.org/clustal2/). The numbers of amino acids are marked on the right. The column below shows the conservation
among different species. The dot ('.'), conservative; ' $\because$ ', obviously conservative; '*', absolutely conservative. The red frame shows the 370th amino acid of TOP1 $\alpha$ in Arabidopsis thaliana and conserved amino acids among different species. (B) Phylogenetic tree and genetic distance of Topoisomerase 1 protein sequences among different species. The numbers above indicate the genetic distance.

Table S1. Oligonucleotides used in this study

## Primers for ChIP and qRT-PCR

| Number | Name | Forward (F) and reverse (R) primers |
| :---: | :---: | :---: |
| H3111 | $F L C 1(-662$ to -412$)$ | F: 5'-AGGCGAGTGGTTCTTTGTTTT-3' |
| H3112 |  | R: 5'-CCTCCCCTACGATACGGATT-3' |
| H6342 | $F L C 2$ (-269 to -201) | F: 5'-CTCGTCATGCGGTACACGTGGC-3' |
| H6461 |  | R: 5'-AAAAACCAAATATGTGAATAAAAAC-3' |
| H5744 | $F L C 3(-200$ to -1$)$ | F: 5'-TTGCATCACTCTCGTTTACCC-3' |
| H5745 |  | R: 5'-GGCTTCTCTCCGAGAGGGC-3' |
| H1248 | $F L C 4(+71$ to +150$)$ | F: 5'-TCGCAACGGTCTCATCGA-3' |
| H1249 |  | R: 5'-GGCGGAGACGACGAGAAG-3' |
| H3115 | $F L C 5(+201$ to +388$)$ | F: 5'-ACCTGGGTTTTCATTTGTTCC-3' |
| H3116 |  | R: 5'-TTTGGTTATCTCATGTATCTATC-3' |
| H3117 | $F L C 6(+688$ to +865$)$ | F: 5'-TCATTGGATCTCTCGGATTTG-3' |
| H3118 |  | R: 5'-ACTAATTTGGATAATCACCAAG-3' |
| H3119 | $F L C 7(+1306$ to +1524$)$ | F: 5'-TTCCCACTCTTGCAGTTACACACA-3' |
| H3120 |  | R: 5'-AAGACACAAGATACAAAGGTTGT-3' |
| H3121 | $F L C 8(+2562$ to +1746$)$ | F: 5'-TGAACTCATGAAAGAGGCGTT-3' |
| H3122 |  | R: 5'-TACAAAGCGTGTTATCAAAACC-3' |
| H6339 | $F L C 9(+5506$ to +5616$)$ | F: 5'-ATGGAGAATAATCATCATGTG-3' |
| H6340 |  | R: 5'-CTAATTAAGTAGTGGGAGAG-3' |
| H3123 | $F L C 10(+5733$ to +5869$)$ | F:5'-GTTTGTATATCTTAATACTCTCTCTTTGGC-3' |
| H3124 |  | R: 5'-ATGCAATTCTCACACGAATAAG-3' |
| H2660 | $F P A$ | F: 5'-ACCAAGCACTACGATTGCAGC-3' |
| H2661 |  | R: 5'-ACCTGAAGACTGTTGCTGCTG-3' |
| H2662 | $F Y$ | F: 5'-TCAAGGACAACCAAACAGTG-3' |
| H2663 |  | R: 5'-TGCCTACTGATGTTGCTGATTG-3' |
| H2664 | $F C A$ | F: 5'-AGCAGCAACCGCTACAAAAGATG-3' |


| H2665 |  | R: 5'-TGCGAGAACTGGCACAAAC-3' |
| :---: | :---: | :---: |
| H2821 | FVE | F: 5'-TGCGAGAACTGGCACAAAC-3' |
| H2822 |  | R: 5'-AGGCGAACCAACTCCATTAG-3' |
| H2823 | FLK | F: 5'-ACGTCGGGTTCAAACATAAG-3' |
| H2824 |  | R: 5'-TTGCTGCTCTGGTGCTAC-3' |
| H2825 | $F L D$ | F: 5'-ACGCAGTGACTCGTGTTC-3' |
| H2826 |  | R: 5'-AGGGTATCGCCTTGTTG-3' |
| H2827 | $L D$ | F: 5'-TCGTCACAGGGTCCAAAAC-3' |
| H2828 |  | R: 5'-TATAAAGGGCACGCATC-3 |
| H2670 | VRN2 | F: 5'-ATGGACTTGTCGACTCAGCCAC-3' |
| H2671 |  | R: 5'-TGTCATTCGGATGATCCACAATG-3' |
| H2672 | VIP4 | F: 5'-TGAAGAAGAGGAAGAGGTTGC-3' |
| H2673 |  | R: 5'-TCGTCACTGTCATCAATCACG-3' |
| H2656 | FRI | F: 5'-TGACTGAAGGAGGATTAGCTG-3' |
| H2657 |  | R: 5'-TCTCATTCGAACCACTCATC-3' |
| H1248 | FLC | F: ''-TCGCAACGGTCTCATCGA-3' $^{\prime}$ |
| H1249 |  | R: $5^{\prime}$-GGCGGAGACGACGAGAAG-3' |
| H0999 | $F T$ | F: 5'-TACGAAAATCCAAGTCCCACTG-3' |
| H1000 |  | R: 5'-AAACTCGCGAGTGTTGAAGTTC-3' |
| H0504 | SOC1 | F: 5'-AGGAACATGCTCAATCGAGG-3' |
| H0505 |  | R: 5'-CTTATACACTCTCAGTACTGC-3' |
| H5028 | SDG8 | F: 5'-ACCTGACTTACTCCAATGAGATC-3' |
| H0723 |  | R: 5'-TTAACTGTTGAGCTTCTTCTCTAAA-3' |
| H6529 | CTD | F: 5'-AGCAAGCCCAGACTACAGC-3' |
| H6530 |  | R: 5'-CAGGGTTGCCTTTATCATCC-3' |
| H0069 | Tublin | F: 5'-GAGCCTTACAAGCTACTCTGTCTGTC-3' |
| H0070 |  | R: 5'-ACACCAGACATAGTAGCAGAAATCAAG-3' |

Primers for in situ hybridization

| Number | Name | Forward (F) and reverse (R) primers |
| :--- | :--- | :--- |
| H0079 | TOP1 $\alpha$ | T7: 5'-TAATACGACTCACTATAGGG-3' |
| H0080 |  | Sp6: 5'-ATTTAGGTGACACTATAGAATACT-3' |

## Primers for cloning

| Number | Name | Forward (F) and reverse (R) primers |
| :---: | :---: | :---: |
| H5603 | SDG8 | F: 5'-GCGTCGACCTAAAAAACCATGTTGGGGATTCA-3' |
| H5604 |  | R: 5'-TTTTCCTTTTGCGGCCGCAAATTTAACTTTCAAACGAAGGC-3' |
| H5308 | TOP $1 \alpha-1$ | F : |
| H4036 |  | 5'-ATAAGAATGCGGCCGCCATGGGCACTGAAACAGTTTCAAAACC-3' |
|  |  | R: 5'-CCGCTCGAGTTATTTCTTTTGCCCATCTCCAGAGGAAG-3' |
| H5309 | TOP1 1 -2 | F : |
| H4038 |  | 5'-ATAAGAATGCGGCCGCcAAATGGACTACTTTGGTGCACAACGG-3' |
|  |  | R: 5'-CCGCTCGAGTTATCCCAAAAATACATACTTGAATTC-3' |
| H5310 | TOP1 $\alpha-3$ | F : |
| H4041 |  | 5'-ATAAGAATGCGGCCGCCCTTTTCAGAGGCCGTGGAGAACATCC-3' |
|  |  | R: 5'-ACGCGTCGACACACACATGGTGCGCAAATTGAAAAATTG-3' |
| H5579 | CTD | F: 5'-ACGCGTCGACaGTTTATCCCCAATGTCAGATGCAC-3' |
| H5580 |  | R: 5'-ATAGTTTAGCGGCCGCCAGGGTTGCCTTTATCATCCTTAC-3' |
| H3109 | FLC | F: 5'-ACGCGTCGACATGGGAAGAAAAAAACTAGAAATC-3' |
| H3110 |  | R: 5'-TTTTCCTTTTGCGGCCGCCTAATTAAGTAGTGGGAGAG-3' |

