

This product is for research use only (not for diagnostic or therapeutic use)

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Product no AS06 198

## Anti-FT/TSF | Flowering locus T and twin sister of FT

## **Product information**

Immunogen KLH-conjugated synthetic peptide derived from A. thaliana FT protein sequence (Q9SXZ2, At1g65480); please note that this antibody will cross-react with the highly homologous TSF (twin sister of FT) protein

**Host** Rabbit

Clonality Polyclonal

**Purity** Immunogen affinity purified serum in PBS pH 7.4.

Format Lyophilized

Quantity 50 ug

**Reconstitution** For reconstitution add 50 μl of sterile water

Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please Storage remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to

the cap or sides of the tube.

## Application information

Recommended dilution 1:1000 (WB)

Expected | apparent

MW

20 | 20 kDa (Arabidopsis thaliana)

Predicted reactivity Betula luminifera, Brassica napus, Brassica oleracea, Brassica rapa, Citrus sinensis, Eucalyptus sp., Hordeum vulgare, Jatropha curcas, Nicotiana tabacum, Persea americana var. americana, Populus tomentosa, Prunus armeniaca,

Prunus avium, Prunus dulcis, Prunus mume, Prunus persica, Solanum tuberosum, Zea mays, Vitis vinifera

Species of your interest not listed? Contact us

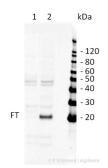
Not reactive in Medicago truncatula, Pisum sativum

Additional information Note that detection for this product is limited by target threshold level

Selected references

Nakamura et al. (2019). High-Resolution Crystal Structure of Arabidopsis FLOWERING LOCUS T Illuminates Its Phospholipid-Binding Site in Flowering. iScience. 2019 Nov 22;21:577-586. doi: 10.1016/j.isci.2019.10.045. Liang and Ow et al. (2019). Nucleocytoplasmic OXIDATIVE STRESS 2 can relocate FLOWERING LOCUS T. Biochemical and Biophysical Research Communications Volume 517, Issue 4, 1 October 2019, Pages 735-740

## Application example



35 μg of total leaf protein extracted with PEB (AS08 300) from wt Arabidopsis thaliana (1) and Arabidopsis thaliana transformed with 35S::FT (2) were separated on 4-12% NuPage (Invitrogen) LDS-PAGE and blotted 80 min (30V) to PVDF. Filter was blocked 1h with 2% low-fat milk powder in TBS-T (0.1% TWEEN 20) and probed with anti-FT/TFT (AS06 198, 1:1000, 1h) and secondary anti-rabbit (1:20 000, 1h) antibody (HRP conjugated) in TBS-T containing 2% low fat milk powder. Antibody incubations were followed by washings in TBS-T (15, +5, +5, +5 min). All steps were performed at RT with agitation. Signal was detected with chemiluminescent detection reagent, using a Fuji LAS-3000 CCD (300s, high sensitivity). Please note that this antibody will not detect FT at 35 µg protein loading in the wt leaf material tested.

Arabidopsis thaliana plants were 4 weeks old, grown @ 8 h light with 130-150 μE light @22°C and 16 h dark @18°C.