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Product no AS20 4430

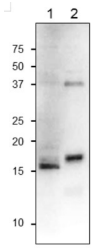
FdC1 | Ferredoxinx-C1

Product information

Immunogen	Purified full length, tag cleaved, recombinant *Arabidopsis thaliana* Ferredoxin C-1, UniProt: O23344 , TAIR: At4g14890
Host	Rabbit
Clonality	Polyclonal
Purity	Total IgG, purified on Protein A
Format	Liquid at 2 mg/ml in PBS, 50% glycerol. Filter sterilized. No preservative or carrier added.
Quantity	100 µg
Storage	Store at -20°C; once make aliquots to avoid repeated freeze-thaw cycles. Please, remember to spin tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tubes.

Application information

Recommended dilution	1: 1000 - 1: 5000 (WB)
Expected | apparent MW	16.7 kDa
Confirmed reactivity	*Arabidopsis thaliana, Zea mays*
Predicted reactivity	*Brassica rapa, Cannabis sativa, Theobroma cacao* Species of your interest not listed? Contact us
Not reactive in	No confirmed exceptions from predicted reactivity are currently known.
Selected references	Voss et al. (2011). FdC1, a Novel Ferredoxin Protein Capable of Alternative Electron Partitioning, Increases in Conditions of Acceptor Limitation at Photosystem I. J Biol Chem. 2011 Jan 7;286(1):50-9. doi: 10.1074/jbc.M110.161562.



10 µg of *Arabidopsis thaliana* total leaf extract **(1)**, 10 µg of *Zea mays* total leaf extract **(2)** were freshly extracted with 2x SDS-sample buffer (+ 2ME) for SDS-PAGE. For IP, 150mM NaCL, 1% Triton X-100, 50 mM Tris-HCl (pH 8.0) and denatured with 4X SDS buffer at 95°C for 5 min. Samples were separated on 10% SDS-PAGE and blotted 1h to PVDF membrane. Blot was blocked with 3 % skim milk/TBS-T, 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 1000 in TBS-T for 1h/RT. The antibody solution was decanted and the blot was washed 4 times for 10 min in TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:10 000 in for 1h/RT with agitation. The blot was washed as above and developed with a chemiluminescent detection reagent, following manufacture's recommendation.