

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

electron microscopy, Western blot)

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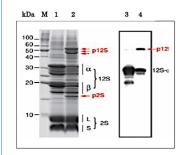
Product no AS20 4403 Anti-12S seed storage protein CRC

Product information

Immunogen	Native, 12S globulin alpaha subunit purified from <i>Arabidopsis thaliana</i> and excised from SDS-PAGE gel. UniProt: <u>Q96318</u> , TAIR: <u>At4g28520</u>	
Host	Rabbit	
Clonality	Polyclonal	
Purity	Total IgG. Protein A purified in PBS, 50% glycerol. Filter sterilized.	
Format	Liquid at 2 mg/ml.	
Quantity	200 µg	
Storage	Store at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please 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.	
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Application information

	Assay dependent (ELISA), 1: 50 (IL by electron microscopy), 1: 100 (IHC), 1: 3000 - 1: 10 000 (WB)
Expected apparent MW	58 30 kDa (alpha subunit)
Confirmed reactivity	Arabidopsis thaliana
Predicted reactivity	Species of your interest not listed? Contact us
Not reactive in	No confirmed exceptions from predicted reactivity are currently known
Selected references	<u>Shirakawa</u> et al. (2014). CONTINUOUS VASCULAR RING (COV1) is a trans-Golgi network-localized membrane protein required for Golgi morphology and vacuolar protein sorting. Plant Cell Physiol. 2014 Apr;55(4):764-72.doi: 10.1093/pcp/pct195. (Immunohistochemistry, Western blot) <u>Li</u> et al. MAG2 and three MAG2-INTERACTING PROTEINs form an ER-localized complex to facilitate storage protein transport in Arabidopsis thaliana. Plant J. 2013 Dec;76(5):781-91.doi: 10.1111/tpj.12347. (Immunolocalisation by



Arabidopsis thaliana dry seed extract from wilde-type (1) and *atvst1-1* mutant (deffective in storage protein transport (2) (SDS-PAGE, stained gel) Western blot of the same samples (3,4) which were separated on 15-20 % SDS-PAGE and blotted to PVDF membrane in wet system. 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: 5000 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 recommendations. P12S is the precursor of 12S globulin. 12S-alpha is the alpha-subunit (proteolyticall form of 12S) of 12S globulin.