# Agrisera

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### Product no AS20 4403

## 12S seed storage protein CRC

#### **Product information**

Immunogen Native, 12S globulin alpaha subunit purified from *Arabidopsis thaliana* and excised from SDS-PAGE gel. UniProt:

Q96318, TAIR: At4g28520

Host Rabbit

Clonality Polyclonal

Purity Total IgG, purified on Protein A/G

Format Liquid at 2 mg/ml in PBS, 50% glycerol. Filter sterilized. No preservative or carrier protein.

Quantity 200 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 assay dependent (ELISA), 1: 50 (IL by electron microscopy), 1: 100 (IHC), 1: 3000 - 1: 10 000 (WB)

Expected | apparent

AW.

58 | 30 kDa (alpha subunit)

Confirmed reactivity

Arabidopsis thaliana

Predicted reactivity

Species of your interest not listed? Contact us

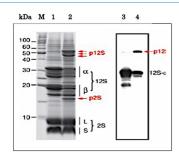
Not reactive in

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)

Li 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 electron microscopy, Western blot)



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.