

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

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## Product no AS20 4405

## Anti-2S3P | 2S seed storage protein 3

## **Product information**

Immunogen Conjugated peptide, derived from N-propeptide of Arabidopsis thaliana 2S3P, UniProt: P15459, TAIR: At4g27160

**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

## **Application information**

Recommended dilution | Assay dependent (ELISA), 1: 500 (IL), 1: 2500 (WB)

Expected | apparent 18.7 | 17-20 kDa (2S albumin precursors)

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

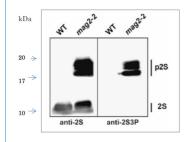
Additional information Anti-2S3P antibodies are recognizing 2S albumin precursors (p2S) but not the mature forms (2S)

Shirakawa et al. (2014). CONTINUOUS VASCULAR RING (COV1) is a trans-Golgi network-localized membrane Selected references

protein required for Golgi morphology and vacuolar protein sorting. Plant Cell Physiol. 2014 Apr;55(4):764-72. doi: 10.1093/pcp/pct195. (Western blot, Arabidopsis thaliana)

Li et al. (2006). MAIGO2 is involved in exit of seed storage proteins from the endoplasmic reticulum in Arabidopsis thaliana. Plant Cell. 2006 Dec;18(12):3535-47. doi: 10.1105/tpc.106.046151.( Immunolocalisation by electron

microscopy, Western blot, Arabidopsis thaliana)



Arabidopsis thaliana dry seeds from wilde-type (WT) and mag2-2 mutant (defective in processing of the precursors) were homogenised in SDS sample buffer (100 mM Tris/HCl, pH 6.8, 4 % SDS w/v, 20 % glycerol v/v, 10 % 2-mercaptoethanol) and separated on 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.

Anti-2S3P antibodies are recognizing 2S albumin precursors (p2S) but not the mature forms (2S).