

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

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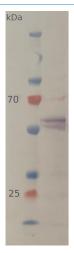
Product no AS23 4916 Anti-UGPase | UDP-glucose pyrophosphorylase (cytoplasm marker, dicots)

Product information

Immunogen	<u>KLH</u> -conjugated peptide derived from <i>Arabidopsis thaliana</i> UGP1, UniProt: <u>A0A1I9LT02</u> and UGP2, UniProt: <u>Q9M9P3</u> TAIR: <u>AT3G03250</u> , <u>AT5G17310</u>
Host	Rabbit
Clonality	Polyclonal
Purity	Antigen affinity purified serum, in PBS pH 7.4
Format	Lyophilized
Quantity	50 µg
Reconstitution	For reconstitution add 50 μ l, of sterile or deionized water.
Storage	Store lyophilized/reconstituted at -20°C; once reconstituted 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 lyophilized material adhering to the cap or sides of the tubes.

Application information

Recommended dilution	1 : 1000 - 1: 5000 (WB)
Expected apparent MW	51.7 52.3 kDa
Confirmed reactivity	Arabidopsis thaliana, Hordeum vulgare, Zea mays
Predicted reactivity	 Arachis hypogaea, Brachypodium distachyon, Brassica napus Cannabis sativa, Capsicum annuum, Glycine max, Gossypium sp., Malus domesticam, Manihot esculenta, Medicago truncatula, Nicotiana tabacum, Pisum sativum, Populus sp., Ricinus communis, Saccharum sp., Solanum lycopresicum, Solanum tuberosum, Sorghum bicolor, Theobroma cacao Species of your interest not listed? <u>Contact us</u> For monocotyl species, use <u>AS14 2813</u>
Not reactive in	Marchantia polymorpha
Selected references	To be added when available, antibody available in March 2024.



10 µg/well of total protein extracted from *Arabidopsis thaliana* leaf, using PEB extraction buffer (Agrisera <u>AS08 300</u>) and denatured with Invitrogen LDS sample buffer (4X) at 70 ° C/5 min. Samples were separated on Invitrogen NuPage Bis-Tris 4-12% SDS-PAGE and blotted for 1 h to Invitrogen PVDF (pore size of 0.45 µm), using: wet transfer. Blot was blocked with 5% milk in TBS-T for: 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 1 000 for 1h/RT with TBS-T Blocking. The antibody solution was decanted, and the blot was rinsed briefly twice, then washed once for 15 min and 3 times for 5 min in TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG ALP conjugated, <u>AS09 607</u> lot 2302) diluted to 1: 5 000 in TBS-T Blocking for 0,5h/RT with agitation. The blot was washed as above and developed with <u>AS19 BCIP-NBT-PLUS</u> lot 08225181 for 1 min. As soon as the desired band is detectable, briefly wash the membrane in generous amounts of deionized water. Transfer the membrane to fresh deionized water and incubate for 2 minutes with agitation before placing



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the membrane on Whatman paper to dry.

Courtesy Agrisera

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