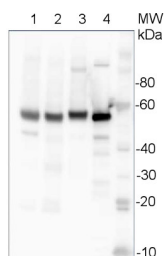


Product no **AS14 2813****Anti-UGPase | UDP-glucose pyrophosphorylase (cytoplasm marker)****Product information**

<b>Immunogen</b>	His-tagged, full length <i>Hordeum vulgare</i> UGPase, overexpressed and purified from <i>E.coli</i> , UniProt: <a href="#">Q43772.1</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Serum
<b>Format</b>	Lyophilized
<b>Quantity</b>	50 µl
<b>Reconstitution</b>	For reconstitution add 50 µl of sterile water
<b>Storage</b>	Store lyophilized/reconstituted 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.
<b>Additional information</b>	Cellular [compartment marker] of cytoplasm, UGPase is a cytoplasmic protein Martz et al, (2002)

**Application information**

<b>Recommended dilution</b>	1 : 10 000 (WB)
<b>Expected   apparent MW</b>	52 kDa
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i> , <i>Hordeum vulgare</i> , <i>Zea mays</i>
<b>Predicted reactivity</b>	<i>Bambusa oldhamii</i> , <i>Brassica pekinensis</i> , <i>Brassica rapa</i> , <i>Capsicum annuum</i> , <i>Cucumis sativus</i> , <i>Dendrobium catenatum</i> , <i>Dendrocalamus sinicus</i> , <i>Glycine max</i> , <i>Gossypium hirsutum</i> , <i>Lycopersicum esculentum</i> , <i>Lycopersicum chilense</i> , <i>Marchantia polymorpha</i> , <i>Oryza sativa</i> , <i>Picea glauca</i> , <i>Populus sp.</i> , <i>Solanum tuberosum</i> , <i>Populus tremula</i> , <i>Ricinus communis</i> , <i>Saccharum officinarum</i> , <i>Vitis vinifera</i> , for more species, please Species of your interest not listed? <a href="#">inquire</a>
	Species of your interest not listed? <a href="#">Contact us</a>
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known
<b>Additional information</b>	This antibody is also recognizing recombinant UGPase, below 0,5 pmol
<b>Selected references</b>	<a href="#">Martín-Merchán</a> et al. (2024). Arabidopsis AGO1 N-terminal extension acts as an essential hub for PRMT5 interaction and post-translational modifications. <i>Nucleic Acids Res.</i> 2024 May 20:gkae387.doi: 10.1093/nar/gkae387. <a href="#">Kleczkowski</a> LA & Decker DD (2015) Sugar activation for production of nucleotide sugars as substrates for glycosyltransferases in plants. <i>J. Appl. Glycosci.</i> (in press).

**application example**

10 µg of total protein from *Arabidopsis thaliana* leaf (1), *Hordeum vulgare* leaf (2), *Zea mays* leaf (3), recombinant UGPase 0.5 pmol (4), were extracted with Protein Extraction Buffer PEB ([AS08 300](#)). Samples were diluted with 1X sample buffer (NuPAGE LDS sample buffer (Invitrogen) supplemented with 50 mM DTT and heat at 70°C for 5 min and kept on ice before loading. Protein samples were separated on 4-12% Bolt Plus gels, LDS-PAGE and blotted for 70 minutes to PVDF using tank transfer. Blots were blocked immediately following transfer in 2% blocking reagent (GE RPN 2125; Healthcare) or 5% non-fat milk dissolved in 20 mM Tris, 137 mM sodium chloride pH 7.6 with 0.1% (v/v) Tween-20 (TBS-T) for 1h at room temperature with agitation. Blots were incubated in the primary antibody at a dilution of 1: 10 000 (in blocking reagent) for 1h at room temperature with agitation. The antibody solution was decanted and the blot was rinsed briefly twice, and then washed 1x15 min and 3x5 min with TBS-T at room temperature with agitation. Blots were incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, recommended secondary antibody [AS09 602](#), Agrisera) diluted to 1:25 000 in blocking reagent for 1h at room temperature with agitation. The blots were washed as above. The blot was developed for 5 min with detection reagent according the manufacturers instructions.

Images of the blots were obtained using a CCD imager (VersaDoc MP 4000) and Quantity One software (Bio-Rad). Exposure time was 10 seconds.