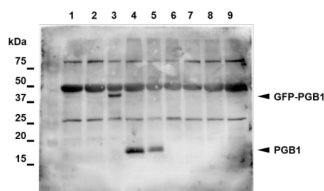


Product no **AS18 4230****AHB1 | Non-symbiotic hemoglobin 1 (A. thaliana)****Product information**

<b>Immunogen</b>	KLH-conjugated peptide derived from <i>Arabidopsis thaliana</i> AHB1, UniProt: <a href="#">O24520</a> , TAIR: <a href="#">AT2G16060</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Antigen affinity purified serum, in PBS pH 7.4.
<b>Format</b>	Lyophilized
<b>Quantity</b>	50 µg
<b>Reconstitution</b>	For reconstitution add 50 µl, of sterile water.
<b>Storage</b>	Lyophilized antibody can be stored at -20°C for up to 3 years. 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**

<b>Recommended dilution</b>	1 : 1000 (WB)
<b>Expected   apparent MW</b>	kDa
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known
<b>Additional information</b>	Conditions for detection of endogenous AHB1 remain to be determined.
<b>Selected references</b>	To be added when available. Antibody released in April 2022.



90 µg/well of total protein extracted freshly from *Arabidopsis thaliana* seedlings with extraction buffer (100mM Tris-HCl, 150mM NaCl, 0.25% NP-40) containing 1mM PMSF and 1X cOmplete® EDTA-free proteases inhibitors, Sigma) and denatured with buffer (Tris-HCl 0,25 M, pH 6,8; SDS 8% (w/v); glycerol 40% (v/v); -Mercaptoethanol 20% (v/v) and Bromophenol Blue) at 90°C for 10 min were separated on 12% SDS-PAGE and blotted (20min; 2A; 25V) to PVDF (pore size of 0.2 µm), using semi-dry transfer. Blot was blocked with 5% milk for 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 1000 ON/4°C with agitation. The antibody solution was decanted and the blot was washed 2 times for 5 min in TBS-T at RT with agitation. Blot was incubated in Agrisera matching secondary antibody (Goat anti-rabbit IgG HRP peroxidase conjugated, [AS09 602](#)) diluted to 1:25 000 in for 1h/RT with agitation. The blot was washed 4 times for 5 min in TBS-T at RT with agitation and developed with Agrisera ECL SuperBright, [AS16 ECL-S](#). Exposure time was 5 seconds.

The lines used in the experiments were generated by:  
 -[Hebeslstrup et al., 2008](#) corresponds to 35S: GFP-PGB1  
 -[Perazzolli et al., 2004](#) corresponds to 35S:PGB1

Courtesy of Dr. Inmaculada Sánchez-Vicente and Isabel Manrique-Gil, USAL-CIALE Salamanca, Spain