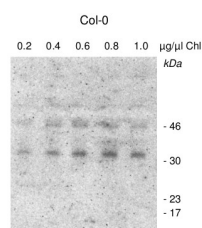


Product no **AS12 1853****Anti-CGL160 | Conserved in green lineage 160****Product information**

<b>Immunogen</b>	part of <i>Arabidopsis thaliana</i> recombinant CGL160 derived from a following sequence UniProt: <a href="#">Q82279</a> , TAIR: <a href="#">At2g31040</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.

**Application information**

<b>Recommended dilution</b>	1 : 5000 (WB)
<b>Expected   apparent MW</b>	38,6   34 kDa (without transit peptide)
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known
<b>Selected references</b>	<a href="#">Galvis</a> et al. (2020). H <sup>+</sup> transport by K <sup>+</sup> EXCHANGE ANTIporter3 promotes photosynthesis and growth in chloroplast ATP synthase mutants. Plant Physiol. pp.01561.2019. doi: 10.1104/pp.19.01561. <a href="#">Fristedt</a> et al. (2015). The Thylakoid Membrane Protein CGL160 Supports CF1CF0 ATP Synthase Accumulation in <i>Arabidopsis thaliana</i> . PLoS One. 2015 Apr 2;10(4):e0121658. doi: 10.1371/journal.pone.0121658.

**application example**

**Respective amounts of** µg/ul of chlorophyll from *Arabidopsis thaliana* leaf extracted with sample buffer (2% SDS, 8% sucrose, 0.2mM EDTA, 10mM Tris HCl (pH 6.8) 4% beta-mercaptoethanol) were separated on 15 % SDS-PAGE and blotted 1h to PVDF. Blots were blocked with 10 % milk for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 4 000 overnight at 4°C with agitation. 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 secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:20 000 in for 1h at RT with agitation. The blot was washed as above and developed for 5 min with ECL according to the manufacturer's instructions. Exposure time was 60 seconds.

Cross-reactivity at around 50 kDa is determined to be CF<sub>1</sub> alpha or beta.

Courtesy of Dr. Rikard Fristedt, Biophysics of Photosynthesis, Dep. Physics and Astronomy, Faculty of Sciences. VU University of Amsterdam, The Netherlands