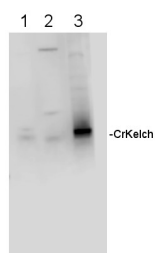


Product no **AS14 2761****Anti-Kelch repeat protein (Chlamydomonas)****Product information**

<b>Immunogen</b>	Recombinant Cr-Klechl, UniProt: <a href="#">A8J2W4</a> , Locus name: g2987 (from <i>Chlamydomonas Reinhardtii</i> )
<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>	1:1000 (WB)
<b>Expected   apparent MW</b>	33.5 kDa
<b>Confirmed reactivity</b>	<i>Chlamydomonas reinhardtii</i> (strain 213grm cw15, M10), <i>Gonium pectorale</i> , <i>Eudorina elegans</i>
<b>Predicted reactivity</b>	Algae
	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

**Application example**

Total protein from *Chlamydomonas reinhardtii* (1), *Gonium Pectorale* (2), *Eudorina elegans* (3) was extracted using 1x Laemmli buffer + DTT + protease and phosphatase inhibitors without bromophenol blue heated at 95 °C for ten minutes then vortexed with zirconia beads at max speed for ten minutes. Lysate was spun at 4 °C for ten minutes at max speed in a tabletop centrifuge. Proteins were precipitated out of supernatant in 80% acetone at -20 °C for 20 min, spun at 4 °C at max speed for 10 min. Pellet was washed 1x with 80% acetone and resuspended in 1x LDS sample buffer + DTT + inhibitors at a concentration of 100 mg chlorophyll/uL. Protein equivalent to 1 µg of chlorophyll was loaded on a 4-20% Bis-Tris gel and blotted to nitrocellulose using wet transfer methods. Blot was blocked with 5% milk in TBS-T for 1hr/RT. Blot was incubated in the primary antibody at a dilution of 1:1000 in 5% milk in TBS-T ON/4°C. The antibody solution was decanted and the blot was rinsed 3 times for 5 min in TBS-T at RT with agitation. Blot was incubated in Agrisera matching secondary diluted to 1:25,000 in 5% milk in TBS-T for 1hr/RT with agitation. The blot was washed as above and developed with chemiluminescent detection reagent and imaged on a Licor C-Digit blot scanner. Exposure time was 12 min.

Courtesy of Jessica Rakijas, Kansas State University, USA