



# SZABO SCANDIC

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## Produktinformation



Forschungsprodukte & Biochemikalien



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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# Agrisera

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

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## Product no **AS16 4083** **HXK1 | Hexokinase 1 (Chlamydomonas)**

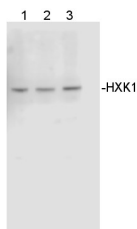
### Product information

<b>Immunogen</b>	KLH-conjugated synthetic peptide derived from <i>Chlamydomonas reinhardtii</i> hexokinase-1, UniProt: <a href="#">A8JGU7</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 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>	67.3 kDa
<b>Confirmed reactivity</b>	<i>Chlamydomonas reinhardtii</i> , <i>Eudorina elegans</i> , <i>Gonium pectorale</i>
<b>Predicted reactivity</b>	Species of your interest not listed? <a href="#">Contact us</a>
<b>Selected references</b>	<a href="#">Upadhyaya</a> and Jagadeeshwar Rao (2019). Reciprocal regulation of photosynthesis and mitochondrial respiration by TOR kinase in <i>Chlamydomonas reinhardtii</i> . Plant Direct Volume 3, Issue 11.

#### Application example



Total protein of *Chlamydomonas reinhardtii* strain 21gr (1), strain cw15 (2), strain M10 (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 1 µg chlorophyll/µL. Protein equivalent to 1 of chlorophyll was loaded on a 4-20% Bis-Tris gel and blotted to PVDF using wet transfer methods. Blot was blocked with 10% milk in TBS-T for 1 hr/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 1:50 000 ([AS09 602](#)) in 5% milk in TBS-T for 1 hr/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