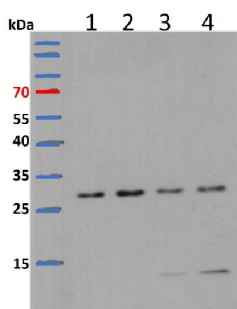


Product no **AS13 2650****Anti-L13-1 | 60S ribosomal protein L13-1****Product information**

<b>Immunogen</b>	KLH-conjugated synthetic peptide derived from <i>Arabidopsis thaliana</i> UniProt: <a href="#">P41127</a> , TAIR: <a href="#">At3g49010</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 steril 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 : 2500 (WB)
<b>Expected   apparent MW</b>	23.7   29 kDa ( <i>Arabidopsis thaliana</i> )
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i> , <i>Hordeum vulgare</i> , <i>Nicotiana benthamiana</i> , <i>Zea mays</i>
<b>Predicted reactivity</b>	<i>Brassica napus</i> , <i>Chlamydomonas reinhardtii</i> , <i>Nicotiana tabacum</i> , <i>Oryza sativa</i> , <i>Picea excelsa</i> , <i>Populus balsamifera</i> , <i>Sorghum bicolor</i> , <i>Ricinus communis</i> , <i>Vitis vinifera</i> Species of your interest not listed? <a href="#">Contact us</a>
<b>Not reactive in</b>	Diatoms
<b>Additional information</b>	Protein or membrane sample should be treated at 70°C for 10 min before loading on the gel
<b>Selected references</b>	<p><a href="#">Wang</a> et al. (2024). Maize Dek51 encodes a DEAD-box RNA helicase essential for pre-rRNA processing and seed development. <i>Cell Rep.</i> 2024 Sep 24;43(9):114673. doi: 10.1016/j.celrep.2024.114673.</p> <p><a href="#">Lee</a> et al. (2023). The Phytophthora nucleolar effector Pi23226 targets host ribosome biogenesis to induce necrotrophic cell death. <i>Plant Commun.</i> 100606. doi: 10.1016/j.xplc.2023.100606.</p> <p><a href="#">Pereira</a> Firmino et al. (2020). Separation and Paired Proteome Profiling of Plant Chloroplast and Cytoplasmic Ribosomes. <i>Plants (Basel)</i> . 2020 Jul 14;9(7):892.doi: 10.3390/plants9070892.</p> <p><a href="#">Shinozaki</a> et al. (2020). Autophagy Increases Zinc Bioavailability to Avoid Light-Mediated ROS Production under Zn Deficiency. <i>Plant Physiol.</i> 2020 Jan 15. pii: pp.01522.2019. doi: 10.1104/pp.19.01522.</p> <p><a href="#">Li</a> (2019). The Isolation of Total and Membrane-Bound Polysomes from Arabidopsis and the Detection of Their Associated AGO1 and sRNAs. <i>Methods Mol Biol.</i> 2019;1932:317-333. doi: 10.1007/978-1-4939-9042-9_23.</p> <p><a href="#">You</a> et al. (2019). FIERY1 promotes microRNA accumulation by suppressing rRNA-derived small interfering RNAs in Arabidopsis. <i>Nat Commun.</i> 2019 Sep 27;10(1):4424. doi: 10.1038/s41467-019-12379-z.</p> <p><a href="#">de Francisco Amorim</a> et al. (2018). The U1 snRNP Subunit LUC7 Modulates Plant Development and Stress Responses via Regulation of Alternative Splicing. <i>Plant Cell.</i> 2018 Nov;30(11):2838-2854. doi: 10.1105/tpc.18.00244. Epub 2018 Oct 11.</p> <p><a href="#">Beine-Golovchuk</a> et al. (2018). Plant Temperature Acclimation and Growth Rely on Cytosolic Ribosome Biogenesis Factor Homologs. <i>Plant Physiol.</i> 2018 Mar;176(3):2251-2276. doi: 10.1104/pp.17.01448. Epub 2018 Jan 30.</p> <p><a href="#">Wang</a> et al. (2016). Comprehensive proteomic analysis of developing protein bodies in maize (<i>Zea mays</i>) endosperm provides novel insights into its biogenesis. <i>J Exp Bot.</i> 2016 Dec;67(22):6323-6335. Epub 2016 Oct 27.</p>



Samples:

- 1 - 15 µg of *Arabidopsis thaliana* whole seedling extract
- 2 - 15 µg of *Arabidopsis thaliana* whole seedling extract
- 3 - 15 µg of *Arabidopsis thaliana* whole seedling treated with Drug-1
- 4 - 15 µg of *Arabidopsis thaliana* whole seedling treated with Drug-2

15 µg/well of total protein extracted freshly from *Arabidopsis thaliana* 8-day-old seedlings extracted with buffer containing 8 M Urea. Samples were separated in the RT on 12 % SDS-PAGE and blotted for 2 h to PVDF (pore size of 0.2 µm), using: wet transfer in the cold. Blot was blocked with 10 % milk for 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 3000 for 1h/RT in PBS 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 PBS at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1: 5000 in for 1 h/RT with agitation. The blot was washed as above and developed with a chemiluminescent detection reagent. Exposure time was 1 minute.

Courtesy of Dr. Kuo-En Chen, Department of Biology, Washington University in St. Louis, USA