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Product no AS15 3030

NodGS | Nodulin / glutamate-ammonia ligase-like protein

Product information

Immunogen KLH-conjugated synthetic peptide derived from a C-terminal of Arabidopsis thaliana NodGS, UniProt: Q8W473,

TAIR: <u>AT3G53180</u>

Host Rabbit

Clonality Polyclonal

Purity Serum

Format Lyophilized

Quantity 50 ul

Reconstitution For reconstitution add 50 μl of sterile water

Storage 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

Recommended dilution 1:300-1:400 (IP), 1:3000 (IF), 1:2000 (WB)

Expected | apparent

MW 94 | 90 kDa

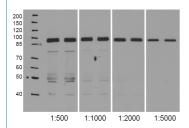
Species of your interest not listed? Contact us

Not reactive in No confirmed exceptions from predicted reactivity are currently known

Selected references Doskocilova et al. (2011). A nodulin/glutamine synthetase-like fusion protein is implicated in the regulation of root

morphogenesis and in signalling triggered by flagellin. Planta 234, 459-476. doi: 10.1007/s00425-011-1419-7

Application example



4,5 of total protein (supernatant after 1h at 25.000 xg) from *Arabidopsis thalian*a suspension culture extracted with 50mM HEPES pH 7,5/75mM NaCl/1mM EGTA/1mM MgCl2/1mM NaF in the presence of protease inhibitors were separated on 8 % SDS-PAGE and blotted 1,5h to a nitrocellulose membrane using tank transfer. Blots were blocked with 5% milk for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 500, 1:1000, 1:2.000, and 1:5000 overnight at 4°C without agitation. The antibody solution was decanted and the blot was rinsed, then washed 3 times for 10 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:10 000 in 5% milk for 30 min at RT with agitation. The blot was washed as above and developed for 1 min with chemiluminescent detection reagent. Exposure time was 15 seconds.

Courtesy of Dr. Jana Chumová, Laboratoř funkční cytologie, Czech Republic