

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

contact: support@agrisera.com

Agrisera AB | Box 57 | SE-91121 Vännäs | Sweden | +46 (0)935 33 000 | www.agrisera.com

Product no AS22 4832

## Anti-Ble tag | Bleomycin resistance protein

## **Product information**

**Immunogen** KLH-conjugated peptide derived from Bleomycin resistance protein from *Streptoalloteichus hindustanus*, UniProt:

P17493

**Host** Rabbit

Clonality Polyclonal

**Purity** Antigen affinity purified serum, in PBS pH 7.4

Format Lyophilized

Quantity 50 μg

**Reconstitution** For reconstitution, add 50 μl of sterile or deionized water.

Storage 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**

**Recommended dilution** 1 : 5000 - 1: 25 000 (WB)

Expected | apparent MW 13.796 kDa

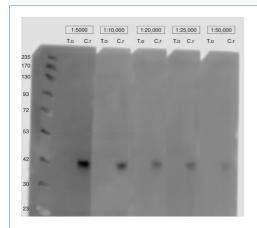
Confirmed reactivity Ble tag

IVIVV

Predicted reactivity | Species of your interest not listed? Contact us

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

Selected references To be added when available, antibody available in July 2024.



2.5 µg/well of total protein extracted freshly from *Tetradesmus obliquus*. Exact buffer components were: PBS and denatured with exact buffer components NuPAGE™ LDS Sample Buffer (2X) (Invitrogen) at 95°C 10 min. Samples were separated in NuPAGE™ Bis-Tris Mini Protein Gels, 4–12%, (Thermoscientific) and blotted for 3 mins to nitrocellulose (pore size of 0.2 µm), using: iblot 2. Blot was blocked with 7.5 % milk TBS-T for: Overnight/18C with agitation. Blot was incubated in the primary antibody at a dilution of 1:5 000, 1:10 000, 1:20 000, 1:25 000, 1:50,000 for 1h/RT with agitation in 7.5% milk TBS-T 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 matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:10 000 in for 1h/RT with agitation. The blot was washed as above and developed with a following chemiluminescent detection reagent. Exposure time was 12 minutes using Li-Cor.