Product Datasheet

RHAMM/CD168 Antibody (PSH02-16) NBP3-32129

Unit Size: 100 ul

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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NBP3-32129

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Images

Western Blot: RHAMM/CD168 Antibody (PSH02-16) [NBP3-32129] - Western blot analysis of RHAMM/CD168 on different lysates with Rabbit anti-RHAMM/CD168 antibody (NBP3-32129) at 1/1,000 dilution.

Lane 1: K-562 cell lysate Lane 2: SW480 cell lysate Lane 3: HepG2 cell lysate Lane 4: Raji cell lysate Lane 5: MCF7 cell lysate Lane 6: SK-Br-3 cell lysate Lane 7: HeLa cell lysate

Lane 8: Jurkat cell lysate

Lysates/proteins at 20 ug/Lane.

Predicted band size: 84 kDa Observed band size: 84 kDa

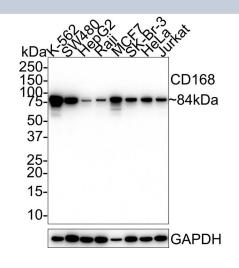
Exposure time: 1 minute 46 seconds;

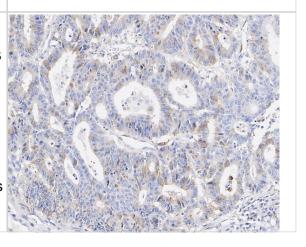
4-20% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (NBP3-32129) at 1/1,000 dilution was used in 5% NFDM/TBST at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody at 1/50,000 dilution was used for 1 hour at room temperature.

Immunohistochemistry: RHAMM/CD168 Antibody (PSH02-16) [NBP3-32129] - Immunohistochemical analysis of paraffin-embedded human colon carcinoma tissue with Rabbit anti-RHAMM/CD168 antibody (NBP3-32129) at 1/1,000 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH2O and PBS, and then probed with the primary antibody (NBP3-32129) at 1/1,000 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.







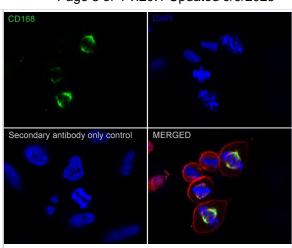
Immunocytochemistry/ Immunofluorescence: RHAMM/CD168 Antibody (PSH02-16) [NBP3-32129] - Immunocytochemistry analysis of HeLa cells labeling RHAMM/CD168 with Rabbit anti-RHAMM/CD168 antibody (NBP3-32129) at 1/100 dilution.

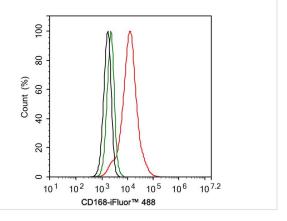
Cells were fixed in 4% paraformaldehyde for 20 minutes at room temperature, permeabilized with 0.1% Triton X-100 in PBS for 5 minutes at room temperature, then blocked with 1% BSA in 10% negative goat serum for 1 hour at room temperature. Cells were then incubated with Rabbit anti-RHAMM/CD168 antibody (NBP3-32129) at 1/100 dilution in 1% BSA in PBST overnight at 4 □. Goat Anti-Rabbit IgG H&L (iFluor™ 488) was used as the secondary antibody at 1/1,000 dilution. PBS instead of the primary antibody was used as the secondary antibody only control. Nuclear DNA was labelled in blue with DAPI.

Beta tubulin (red) was stained at 1/100 dilution overnight at +4□. Goat Anti-Mouse IgG H&L (iFluor™ 594) was used as the secondary antibody at 1/1,000 dilution.

Flow Cytometry: RHAMM/CD168 Antibody (PSH02-16) [NBP3-32129] - Flow cytometric analysis of HeLa cells labeling RHAMM/CD168.

Cells were fixed and permeabilized. Then stained with the primary antibody (NBP3-32129, 1µg/mL) (red) compared with Rabbit IgG Isotype Control (green). After incubation of the primary antibody at +4□ for an hour, the cells were stained with a iFluor™ 488 conjugate-Goat anti-Rabbit IgG Secondary antibody at 1/1,000 dilution for 30 minutes at +4□. Unlabelled sample was used as a control (cells without incubation with primary antibody; black).









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NB7160 Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]

NBP2-24891 Rabbit IgG Isotype Control

NBP2-56490PEP RHAMM/CD168 Recombinant Protein Antigen

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