

Product Data Sheet

Anti-Visfatin Antibody

Catalog # Source Reactivity Applications

CQA8218 Rabbit H, M, R WB, IH, IF/IC, IP

Description Rabbit monoclonal antibody to Visfatin

Immunogen A synthetic peptide of human Visfatin

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Visfatin protein.

Clonality Monoclonal

Conjugation

Form Liquid in 50mM Tris-Glycine (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide

and 0.05% BSA.

Dilution WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/100), IP (1/10 - 1/50)

Gene Symbol NAMPT

Alternative Names PBEF; PBEF1; Nicotinamide phosphoribosyltransferase; NAmPRTase; Nampt;

Pre-B-cell colony-enhancing factor 1; Pre-B cell-enhancing factor; Visfatin

Entrez Gene 10135 (Human); 59027 (Mouse); 297508 (Rat)

SwissProt P43490 (Human); Q99KQ4 (Mouse); Q80Z29 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

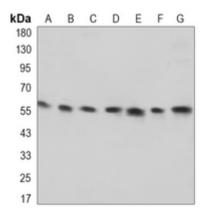
freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

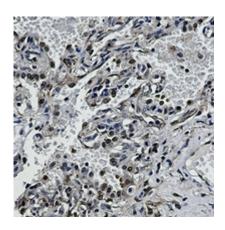
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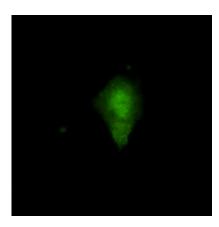
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Western blot analysis of Visfatin expression in NIH3T3 (A), Raw264.7 (B), Hela (C), A549 (D), HL60 (E), U2OS (F), C6 (G) whole cell lysates. (Predicted band size: 56 kD; Observed band size: 56 kD)



Immunohistochemical analysis of Visfatin staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of Visfatin staining in U87MG cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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