

Product Data Sheet

Anti-Visfatin Antibody

Catalog #	Source	Reactivity	Applications
CQA2092	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rabl	oit polyclonal antibody	to Visfatin
Immunogen	Reco	ombinant full length pro	otein of human Visfatin
Purification	The	antibody was purified l	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	els of Visfatin protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium _l	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	NAM	1PT	
Alternative Na	ames PBEI	; PBEF1; Nicotinamide	phosphoribosyltransferase; NAmPRTase; Nampt;
	Pre-	B-cell colony-enhancing	g factor 1; Pre-B cell-enhancing factor; Visfatin
Entrez Gene	1013	35 (Human); 59027 (Mo	ouse); 297508 (Rat)
SwissProt	P434	190 (Human); Q99KQ4	(Mouse); Q80Z29 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid
	free	ze/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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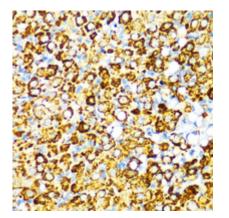
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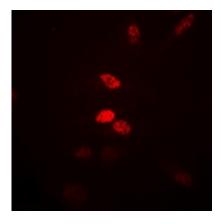
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Western blot analysis of Visfatin expression in mouse liver (A), mouse skeletal muscle (B) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 56 kD)



Immunohistochemical analysis of Visfatin staining in rat ovary 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 Hela 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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