

Product Data Sheet

Anti-Vimentin Antibody

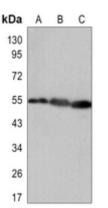
Catalog #	Source	Reactivity	Applications	
CQA8027	Rabbit	H, M, R, Ha	WB, IH, IF/IC	
Description		Rabbit monoclonal antibody	to Vimentin	
Immunogen		A synthetic peptide of huma	n Vimentin	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of Vimentin 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)	
Gene Symbol		VIM		
Alternative Na	ames	Vimentin		
Entrez Gene		7431 (Human); 22352 (Mous	e); 81818 (Rat)	
SwissProt	vissProt P08670 (Human); P20152 (Mouse); P31000 (Rat)			
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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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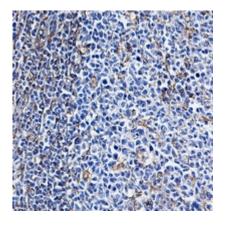
Coherion



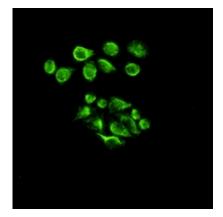
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Western blot analysis of Vimentin expression in C6 (A), CHOK1 (B), Hela (C) whole cell lysates. (Predicted band size: 54 kD; Observed band size: 54 kD)



Immunohistochemical analysis of Vimentin staining in human tonsil 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 Vimentin 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 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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