

Product Datasheet

Vimentin Mouse Monoclonal Antibody (orb1474277)

Catalog Number	orb1474277
Category	Antibodies
Description	The Vimentin Antibody is suitable for IF, IHC, IP, WB. It is a Monoclonal, Unconjugated antibody which raised against Purified recombinant human Vimentin protein fragments expressed in E.coli. Purification: The antibody was purified by immunogen affinity chromatography.
Target	VIM
Clonality	Monoclonal
Species/Host	Mouse
Conjugation	Unconjugated
Reactivity	Human
Form/Appearance	Liquid
Buffer/Preservatives	PBS containing 50% glycerol, 0.5% rAlbumin and 0.02% sodium azide, pH 7.3.
Purification	The antibody was purified by immunogen affinity chromatography.
Immunogen	Purified recombinant human Vimentin protein fragments expressed in E.coli.
UniProt ID	P08670
Tested applications	IF, IHC, IP, WB
Dilution range	WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/100), IP (1/10 - 1/50)

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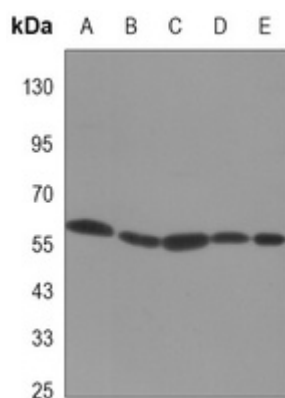
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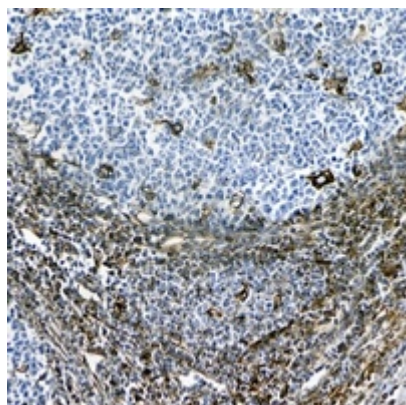
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Specificity	Recognizes endogenous levels of Vimentin protein.
Antibody Type	Primary Antibody
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Note	For research use only
Entrez	7431
Expiration Date	12 months from date of receipt.



Western blot analysis of Vimentin expression in Molt4 (A), K562 (B), COS7 (C), Jurkat (D), HeLa (E) whole cell lysates. (Predicted band size: 54 kD; Observed band size: 57 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 6.136). 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.

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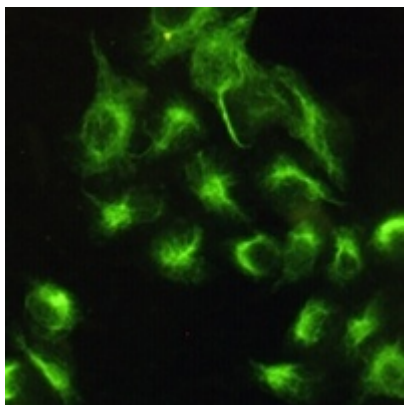
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Immunofluorescent analysis of Vimentin staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified 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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