

### OriGene Technologies, Inc.

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# Product datasheet for TA801117

### Vimentin (VIM) Mouse Monoclonal Antibody [Clone ID: OTI1H10]

### **Product data:**

Product Type:	Primary Antibodies
Clone Name:	OTI1H10
Applications:	IF, WB
Recommended Dilution:	WB 1:200 - 1:1000
Reactivity:	Human, Mouse, Rat
Host:	Mouse
lsotype:	lgG2a
Clonality:	Monoclonal
Immunogen:	Human recombinant protein fragment corresponding to amino acids 210-466 of human VIM (NP_003371) produced in E.coli.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	1 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	53.5 kDa
Gene Name:	vimentin
Database Link:	<u>NP_003371</u> <u>Entrez Gene 22352 MouseEntrez Gene 81818 RatEntrez Gene 7431 Human</u> <u>P08670</u>



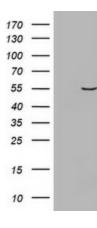
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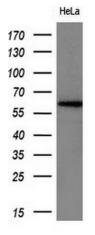
# **Vimentin (VIM) Mouse Monoclonal Antibody [Clone ID: OTI1H10] - TA801117Background:**This gene encodes a member of the intermediate filament family. Intermediate filamentents,<br/>along with microtubules and actin microfilaments, make up the cytoskeleton. The protein<br/>encoded by this gene is responsible for maintaining cell shape, integrity of the cytoplasm, and<br/>stabilizing cytoskeletal interactions. It is also involved in the immune response, and controls<br/>the transport of low-density lipoprotein (LDL)-derived cholesterol from a lysosome to the site<br/>of esterification. It functions as an organizer of a number of critical proteins involved in<br/>attachment, migration, and cell signaling. Mutations in this gene causes a dominant,<br/>pulverulent cataract. [provided by RefSeq, Jun 2009]Synonyms:CTRCT30; HEL113

Protein Families:

## ES Cell Differentiation/IPS

### **Product images:**

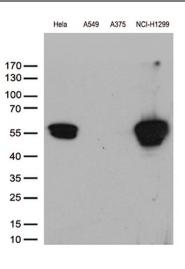


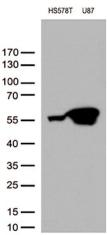


HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY VIM ([RC201546], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-VIM. Positive lysates [LY401165] (100ug) and [LC401165] (20ug) can be purchased separately from OriGene.

Western blot analysis of extracts (10ug) from 1 cell line by using anti-VIM monoclonal antibody at 1:200.

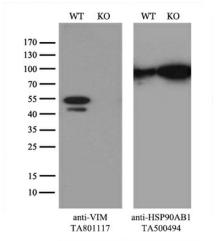
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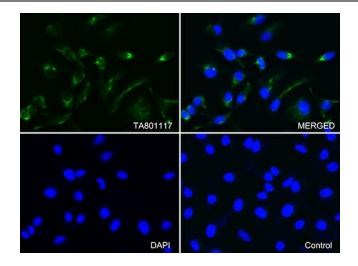


Western blot analysis of extracts (35ug) from 4 different cell lines by using anti-VIM monoclonal antibody (1:500).

Western blot analysis of extracts (35ug) from 2 different cell lines by using anti-VIM monoclonal antibody (1:500).



Equivalent amounts of cell lysates (10 ug per lane) of wild-type Hela cells (WT, Cat# LC810HELA) and VIM-Knockout Hela cells (KO, Cat# [LC810257]) were separated by SDS-PAGE and immunoblotted with anti-VIM monoclonal antibody TA801117, (1:500). Then the blotted membrane was stripped and reprobed with anti-HSP90AB1 antibody ([TA500494]) as a loading control.

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Immunofluorescent staining of Hela cells using anti-VIM mouse monoclonal antibody (TA801117, green, upper left; merged, upper right) or Isotype control (merged, lower right). Cell nuclei were stained with DAPI (blue, lower left) (1:100).

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