

Product datasheet for CF501166

OriGene Technologies, Inc.

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PIM2 Mouse Monoclonal Antibody [Clone ID: OTI9A5]

Product data:

Product Type: Primary Antibodies

Clone Name: OTI9A5

Applications: FC, IF, IHC, IP, WB

Recommended Dilution: WB 1:2000, IHC 1:50, IF 1:100, FLOW 1:100, IP 2ug/500ul

Reactivity: Human, Mouse, Rat

Host: Mouse Isotype: IgG2a

Clonality: Monoclonal

Immunogen: Full length human recombinant protein of human PIM2 (NP_006866) produced in HEK293T

cell

Formulation: Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose)

Reconstitution Method: For reconstitution, we recommend adding 100uL distilled water to a final antibody

concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific)

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: 34.0 kDa

Gene Name: Homo sapiens Pim-2 proto-oncogene, serine/threonine kinase (PIM2), mRNA.

Database Link: NP 006866

Entrez Gene 18715 MouseEntrez Gene 317366 RatEntrez Gene 11040 Human

Q9P1W9

Background: This gene encodes a protooncogene that acts as a serine/threonine protein kinase. Studies

determined the encoded protein functions to prevent apoptosis and to promote cell survival.

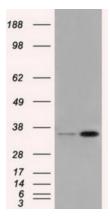




Protein Families: Druggable Genome, Protein Kinase

Protein Pathways: Acute myeloid leukemia

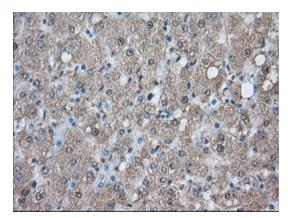
Product images:



HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY PIM2 ([RC201933], 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-PIM2. Positive lysates [LY416355] (100ug) and [LC416355] (20ug) can be purchased separately from OriGene.

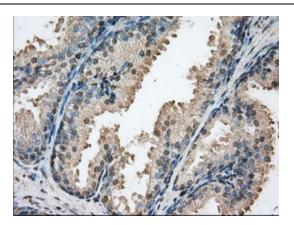


Immunohistochemical staining of paraffinembedded Human colon tissue within the normal limits using anti-PIM2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501166])

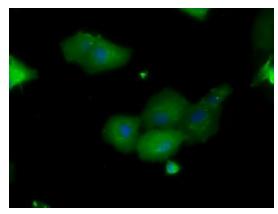


Immunohistochemical staining of paraffinembedded Human liver tissue within the normal limits using anti-PIM2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501166])

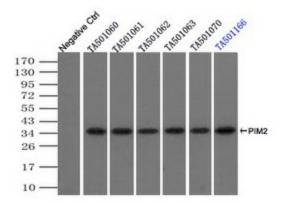




Immunohistochemical staining of paraffinembedded Human prostate tissue within the normal limits using anti-PIM2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501166])

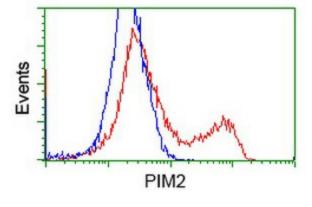


Anti-PIM2 mouse monoclonal antibody ([TA501166]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY PIM2 ([RC201933]).

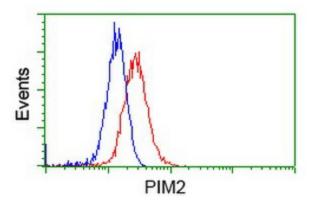


Immunoprecipitation (IP) of PIM2 by using TrueMab monoclonal anti-PIM2 antibodies (Negative control: IP without adding anti-PIM2 antibody.). For each experiment, 500ul of DDK tagged PIM2 overexpression lysates (at 1:5 dilution with HEK293T lysate), 2ug of anti-PIM2 antibody and 20ul (0.1mg) of goat anti-mouse conjugated magnetic beads were mixed and incubated overnight. After extensive wash to remove any non-specific binding, the immunoprecipitated products were analyzed with rabbit anti-DDK polyclonal antibody.

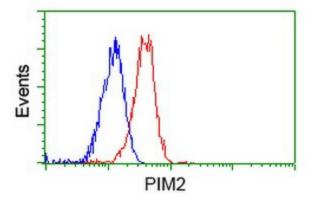




HEK293T cells transfected with either [RC201933] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-PIM2 antibody ([TA501166]), and then analyzed by flow cytometry.



Flow cytometric Analysis of Hela cells, using anti-PIM2 antibody ([TA501166]), (Red), compared to a nonspecific negative control antibody, (Blue).



Flow cytometric Analysis of Jurkat cells, using anti-PIM2 antibody ([TA501166]), (Red), compared to a nonspecific negative control antibody, (Blue).