

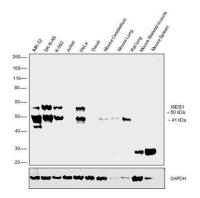


MEIS1 Monoclonal Antibody (OTI2A3)

Product Details	
Size	100 μL
Species Reactivity	Human, Mouse, Rat
Host/Isotype	Mouse / IgG1
Class	Monoclonal
Туре	Antibody
Clone	OTI2A3
Conjugate	Unconjugated
Immunogen	Human recombinant protein fragment corresponding to amino acids 176-390 of MEIS1 produced in E.coli
Form	Liquid
Concentration	1 mg/mL
Purification	Affinity Chromatography
Storage buffer	PBS, pH 7.3, with 1% BSA, 50% glycerol
Contains	0.02% sodium azide
Storage conditions	-20° C, Avoid Freeze/Thaw Cycles
RRID	AB_2724671

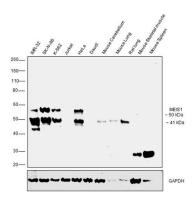
Applications	Tested Dilution	Publications
Western Blot (WB)	1:2,000	-
Immunohistochemistry (Paraffin) (IHC (P))	1:500	-
Immunocytochemistry (ICC/IF)	1:500	-

Product Images For MEIS1 Monoclonal Antibody (OTI2A3)



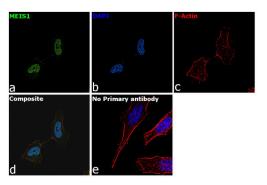
MEIS1 Antibody (MA5-27191)

Antibody specificity was demonstrated by detection of differential basal expression of the target across cell lines and tissues tested owing to their inherent genetic constitution. Relative expression of MEIS1 was observed in IMR-32, SK-N-AS, K-562, HeLa in comparison to Jurkat, Daudi and in Mouse Cerebellum, Mouse Lung, Rat Lung in comparison to Mouse Skeletal Muscle and Mouse Spleen using Anti-MEIS1 Monoclonal Antibody (Product # MA5-27191) in Western Blot. {RE}



MEIS1 Antibody (MA5-27191) in WB

Western blot was performed using Anti-MEIS1 Monoclonal Antibody (Product # MA5-27191). A 50 kDa band and a 41kDa band corresponding to different isoforms of MEIS1 was observed in IMR-32, SK-N-AS, K-562, HeLa cell lines. The 50 kDa band corresponding to MEIS1 was also observed in Mouse Cerebellum, Mouse Lung, Rat Lung and Mouse Skeletal muscle. A noncharacterized band corresponding to 58 kDa was observed in the cell lines tested. Modified Whole cell extracts (30 µg lysate) of IMR-32 (Lane 1), SK-A-NS (Lane 2), K-562 (Lane 3), Jurkat (Lane 4), HeLa (Lane 5), Daudi (Lane 6), tissue extracts of Mouse Cerebellum (Lane 7), Mouse Lung (Lane 8), Rat Lung (Lane 9), Mouse Skeletal Muscle (lane 10) and Mouse Spleen (Lane 11) were electrophoresed using NuPAGE™ 4-12% Bis-Tris Protein Gel (Product # NP0322BOX). Resolved proteins were then transferred onto a nitrocellulose membrane (Product # IB23001) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with the primary antibody (1:500 dilution) and detected by chemiluminescence Goat Anti-Mouse IgG Secondary Antibody, HRP conjugate (Product # A28177, 1:4000 dilution) using the iBright FL 1000 (Product # A32752). Chemiluminescent detection was performed using Novex® ECL Chemiluminescent Substrate Reagent Kit (Product # WP20005).



MEIS1 Antibody (MA5-27191) in ICC/IF

Immunofluorescence analysis of MEIS1 was performed using 70% confluent log phase HeLa cells. The cells were fixed with 4% Paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 15 minutes, and blocked with 2% BSA for 10 minutes at room temperature. The cells were labeled with MEIS1 Monoclonal Antibody (Product # MA5-27191) at 1:500 dilution in 0.1% BSA, incubated at 4 degree Celsius overnight and then labeled with Goat anti-Mouse IgG (H+L) Superclonal™ Secondary Antibody, Alexa Fluor® 488 conjugate (Product # A28175) at a dilution of 1:2000 dilution for 45 minutes at room temperature (Panel a: Green). Nuclei (Panel b: Blue) were stained with ProLong™ Diamond Antifade Mountant with DAPI (Product # P36962). F-actin (Panel c: Red) was stained with Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing Cytoplasmic localization. Panel e represents control cells with no primary antibody to assess background. The images were captured at 60X magnification.

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