

#### Anti-LMO4 Antibody

Rabbit polyclonal antibody to LMO4 Catalog # AP61307

#### Specification

## Anti-LMO4 Antibody - Product Information

Application Primary Accession Other Accession Reactivity Host Clonality Calculated MW WB, IF/IC, IHC <u>P61968</u> <u>P61969</u> Human, Mouse, Rat, Zebrafish, Pig, Bovine Rabbit Polyclonal 17994

#### Anti-LMO4 Antibody - Additional Information

Gene ID 8543

**Other Names** LIM domain transcription factor LMO4; Breast tumor autoantigen; LIM domain only protein 4; LMO-4

**Target/Specificity** KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of human LMO4. The exact sequence is proprietary.

Dilution WB~~WB (1/500 - 1/1000), IH (1/50 - 1/200), IF/IC (1/100 - 1/500) IF/IC~~N/A IHC~~1:100~500

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

## Anti-LMO4 Antibody - Protein Information

Name LMO4

Function

Transcription cofactor. Plays a role in establishing motor neuron identity, in concert with MNX1, acting, at least in part, to disrupt LDB1-LHX3 complexes thereby negatively modulating interneuron genes in motor neurons.

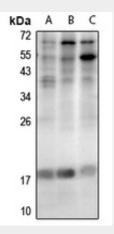


## **Anti-LMO4 Antibody - Protocols**

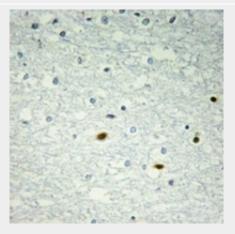
Provided below are standard protocols that you may find useful for product applications.

- <u>Western Blot</u>
- <u>Blocking Peptides</u>
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

# Anti-LMO4 Antibody - Images

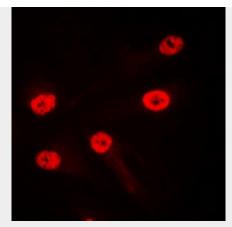


Western blot analysis of LMO4 expression in A549 (A), Hela (B), AML12 (C) whole cell lysates.



Immunohistochemical analysis of LMO4 staining in human brain 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 LMO4 staining in NIH3T3 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 Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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