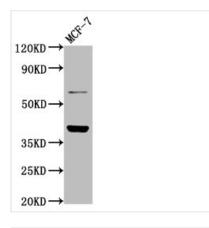






## LAPTM4B Antibody

<b>Product Code</b>	CSB-PA774814LA01HU
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	Q86VI4
Immunogen	Recombinant Human Lysosomal-associated transmembrane protein 4B protein (174-226AA)
Raised In	Rabbit
Species Reactivity	Human
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:500-1:1000, IF:1:50-1:200
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Purification Method	>95%, Protein G purified
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Isotype	IgG
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Isotype	IgG
Isotype Clonality	IgG Polyclonal Lysosomal-associated transmembrane protein 4B (Lysosome-associated
Isotype Clonality Alias	IgG Polyclonal Lysosomal-associated transmembrane protein 4B (Lysosome-associated transmembrane protein 4-beta), LAPTM4B
Isotype Clonality Alias Immunogen Species	IgG  Polyclonal  Lysosomal-associated transmembrane protein 4B (Lysosome-associated transmembrane protein 4-beta), LAPTM4B  Homo sapiens (Human)



Western Blot

Positive WB detected in: MCF-7 whole cell lysate

All lanes: LAPTM4B antibody at 4.8µg/ml

Secondary

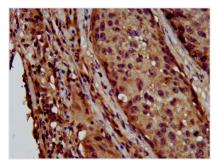
Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 42, 26, 36 kDa Observed band size: 42 kDa

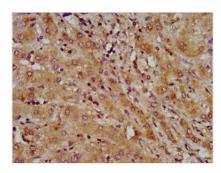




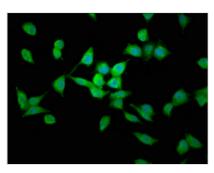




IHC image of CSB-PA774814LA01HU diluted at 1:600 and staining in paraffin-embedded human lung cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



IHC image of CSB-PA774814LA01HU diluted at 1:600 and staining in paraffin-embedded human liver cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence staining of SH-SY5Y cells with CSB-PA774814LA01HU at 1:200, counterstained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).