LAMC2 Antibody



PACO57120

Human

ELISA, WB, IHC

Product Information

Size: Protein Background:

Binding to cells via a high affinity receptor, laminin is thought to mediate the attachment, migration and organization of cells into tissues during embryonic Reactivity:

development by interacting with other extracellular matrix components. Ladsin exerts cell-scattering activity toward a wide variety of cells, including epithelial, endothelial,

and fibroblastic cells.

Source: Gene ID:

Rabbit LAMC2

Isotype: Uniprot

IgG Q13753

Applications: Synonyms:

Recommended dilutions:

ELISA:1:2000-1:10000, WB:1:500-1:5000, IHC:1:200-1:500

Laminin subunit gamma-2 (Cell-scattering factor 140 kDa subunit) (CSF 140 kDa subunit) (Epiligrin subunit gamma) (Kalinin subunit gamma) (Kalinin/nicein/epiligrin 100 kDa subunit) (Ladsin 140 kDa subunit) (Laminin B2t chain) (Laminin-5 subunit gamma) (Large adhesive scatter factor 140 kDa subunit) (Nicein subunit gamma), LAMC2, LAMB2T LAMNB2

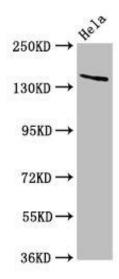
Immunogen:

Recombinant Human Laminin subunit gamma-2 protein (417-588AA).

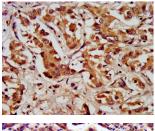
Storage:

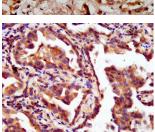
Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

Product Images



Western Blot. Positive WB detected in: Hela whole cell lysate. All lanes: LAMC2 antibody at $5\mu g/ml$. Secondary. Goat polyclonal to rabbit lgG at 1/50000 dilution. Predicted band size: 131, 122 kDa. Observed band size: 140 kDa.





IHC image of PACO57120 diluted at 1:200 and staining in paraffinembedded 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.

IHC image of PACO57120 diluted at 1:200 and staining in paraffinembedded 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.