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# Anti-ADAM12-L Antibody

Catalog # AP53953

## Product Information

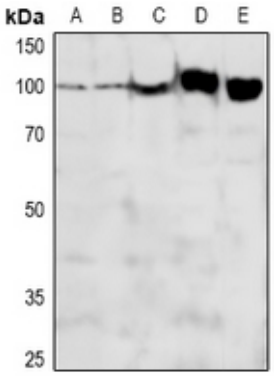
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|---|---|
| **Application** | WB, IHC, IF/IC |
| **Primary Accession** | O43184-1 |
| **Reactivity** | Human, Mouse, Rat |
| **Host** | Rabbit |
| **Clonality** | Polyclonal |

## Additional Information

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|---|---|
| **Other Names** | MLTN; Disintegrin and metalloproteinase domain-containing protein 12; ADAM 12; Meltrin-alpha |
| **Target/Specificity** | KLH-conjugated synthetic peptide encompassing a sequence within the center region of human ADAM12-L. The exact sequence is proprietary. |
| **Dilution** | WB~~1/500 - 1/1000 IHC~~1:100~500 IF/IC~~N/A |
| **Format** | Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide. |
| **Storage** | Store at -20 °C.Stable for 12 months from date of receipt |

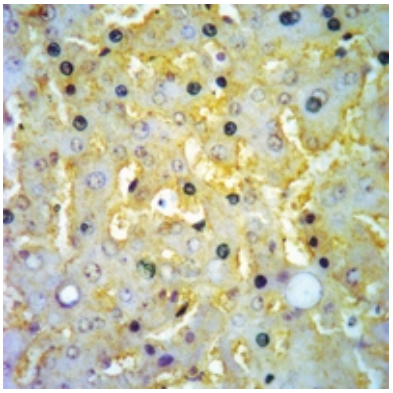
## Protein Information

## Images

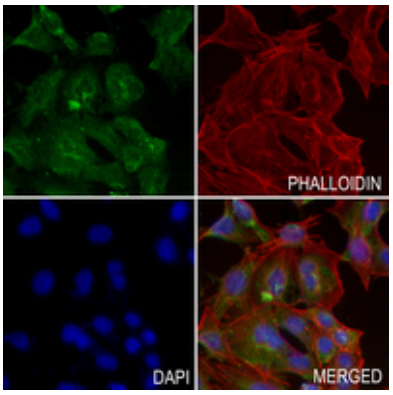



Western blot analysis of ADAM12-L expression in PC3 (A), HCT116 (B), U87MG (C), mouse brain (D), rat brain (E) whole cell lysates. (Predicted band size: 99 kD; Observed band size: 100 kD)

Immunohistochemical analysis of ADAM12-L staining in human liver cancer 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 ADAM12-L staining in MOVAS 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.