## **OCLN Antibody**



## PACO56350

Reactivity:

## **Product Information**

Size: Protein Background:

50ug May play a role in the formation and regulation of the tight junction (TJ) paracellular

permeability barrier. It is able to induce adhesion when expressed in cells lacking tight

junctions.

Human Gene ID:

Source: OCLN

Rabbit Uniprot

**Isotype:** Q16625

lgG Synonyms:

**Applications:** Occludin, OCLN

ELISA, WB, IHC, IF Immunogen:

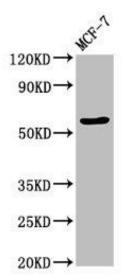
**Recommended dilutions:** Recombinant Human Occludin protein (285-430AA).

ELISA:1:2000-1:10000, WB:1:500-1:5000, IHC:1:200-1:500, IF:1:50-1:200

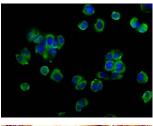
Storage:

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

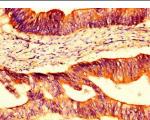
## **Product Images**



Western Blot. Positive WB detected in: MCF-7 whole cell lysate. All lanes: OCLN antibody at  $3.2\mu g/ml$ . Secondary. Goat polyclonal to rabbit lgG at 1/50000 dilution. Predicted band size: 60, 53, 55, 32, 24, 9 kDa. Observed band size: 60 kDa.



Immunofluorescence staining of MCF-7 cells with PACO56350 at 1:133, counter-stained 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).



IHC image of PACO56350 diluted at 1:400 and staining in paraffinembedded human colon 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.