# Occludin Rabbit pAb

Catalog No.: A12621 15 Publications



## **Basic Information**

#### **Observed MW**

62kDa

## **Calculated MW**

59kDa

## Category

Primary antibody

## **Applications**

ELISA,WB

#### **Cross-Reactivity**

Human, Mouse, Rat

## **Background**

This gene encodes an integral membrane protein that is required for cytokine-induced regulation of the tight junction paracellular permeability barrier. Mutations in this gene are thought to be a cause of band-like calcification with simplified gyration and polymicrogyria (BLC-PMG), an autosomal recessive neurologic disorder that is also known as pseudo-TORCH syndrome. Alternative splicing results in multiple transcript variants. A related pseudogene is present 1.5 Mb downstream on the q arm of chromosome 5.

## **Recommended Dilutions**

**WB** 

1:1000 - 1:5000

## **Immunogen Information**

**Gene ID** 100506658

**Swiss Prot** 

Q16625

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 350-450 of human Occludin (NP\_002529.1).

## **Synonyms**

BLCPMG; PTORCH1; PPP1R115; Occludin

## **Contact**

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## **Product Information**

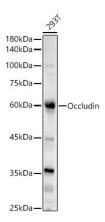
SourceIsotypePurificationRabbitIgGAffinity purification

## Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3.

# **Validation Data**



Western blot analysis of 293T, using Occludin antibody (A12621) at 1:2000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 5s.