

A6827

Leader in Biomolecular Solutions for Life Science



Occludin Rabbit pAb

Catalog No.: A6827

Basic Information

Observed MW

59kDa

Calculated MW

59kDa

Category

Polyclonal Antibody

Applications

WB

Cross-Reactivity

Human, Mouse

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 1:500 - 1:1000

Immunogen Information

Gene ID

100506658

Swiss Prot

Q16625

Immunogen

A synthetic peptide of human Occludin

Synonyms

BLC-PMG; PTORCH1; PPP1R115; Occludin

Contact



www.abclonal.com

Product Information

Source

Rabbit

Isotype

IgG

Purification

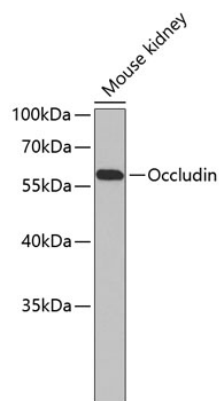
Affinity purification

Storage

Store at 4°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide, pH 7.3.

Validation Data



Western blot analysis of extracts of mouse kidney, using Occludin antibody (A6827).
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.