A2601

Leader in Biomolecular Solutions for Life Science

Occludin Rabbit pAb

Catalog No.: A2601 95 Publications



Basic Information

Observed MW 65kDa/

Calculated MW 59kDa

Category Polyclonal Antibody

Applications WB,ELISA

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

Immunogen Information

WB

1:1000 - 1:5000

www.abclonal.com

Swiss Prot Q16625

Immunogen

Gene ID

100506658

Recombinant fusion protein containing a sequence corresponding to amino acids 400-522 of human Occludin (NP_002529.1).

Synonyms

BLCPMG; PTORCH1; PPP1R115; Occludin

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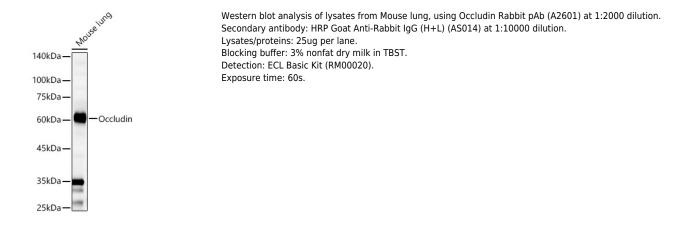
Product Information

Source Rabbit **Isotype** IgG Purification Affinity 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 lysates from Mouse liver using Occludin Rabbit pAb (A2601) at 1:1000 dilution incubated overhight at 4°C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 40 µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 45s. 70kDa—

60kDa —	-	—Occludin
42kDa —		
35kDa —		