

Occludin Rabbit mAb

货号: B31169

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB IP FC
推荐浓度	WB: 1:500 - 1:2000 IP: 1:20 - 1:50 FC: 1:20 - 1:50
理论分子量	8kDa/23kDa/31kDa/52kDa/54kDa/59kDa
实测分子量	59kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	293T,Mouse brain
细胞定位	Cell junction,Membrane,Multi-pass membrane protein,tight junction
纯化	Affinity purification

抗原信息

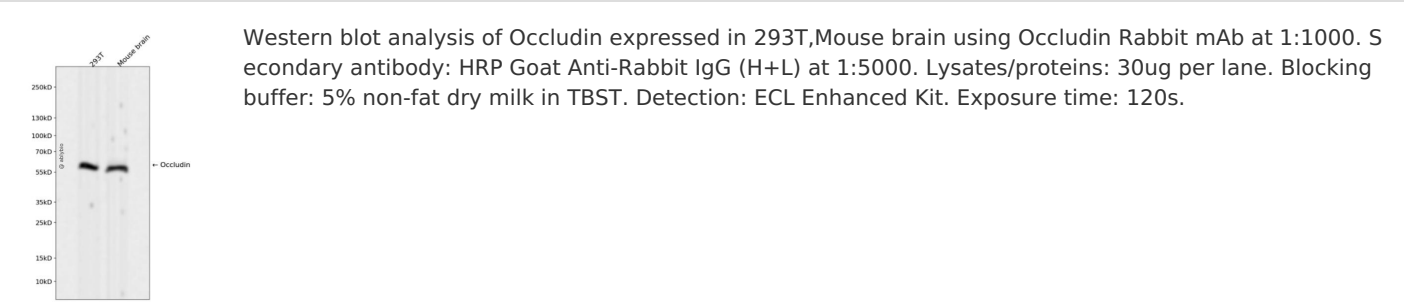
抗原信息	Recombinant fusion protein corresponding to Human Occludin.
序列	TDYTTGGESCDELEEDWIREYPPITSDQQRQLYKRNFDTGLQEYKSLQSELDEINKELSRLDKELDDYREESEEEYMAAADEYNRLKQVKGSADYKSKKNHCKQLKSKLSHIKKMVG DYDRQKT

靶点信息

研究背景	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.
------	---

基因ID	100506658
基因名	OCLN
Swiss	Q16625
别名	OCLN;BLCPMG;PPP1R115;PTORCH1;occludin;Occludin

产品验证



实验步骤

访问官网浏览详情: www.ablybio.cn