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PEX19 (YD14874) Rabbit mAb

货号: **AYD15216**

产品信息

反应	Human,Rat
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB ICC/IF FC IP
推荐浓度	
理论分子量	33kDa
实测分子量	
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	22Rv1,THP-1,U-87MG,Jurkat
细胞定位	Cytoplasm, Peroxisome membrane
纯化	亲和纯化

抗原信息

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靶点信息

研究背景	This gene is necessary for early peroxisomal biogenesis. It acts both as a cytosolic chaperone and as an import receptor for peroxisomal membrane proteins (PMPs). Peroxins (PEXs) are proteins that are essential for the assembly of functional peroxisomes. The peroxisome biogenesis disorders (PBDs) are a group of genetically heterogeneous autosomal recessive, lethal diseases characterized by multiple defects in peroxisome function. These disorders have at least 14 complementation groups, with more than one phenotype being observed for some complementation groups. Although the clinical features of PBD patients vary, cells from all PBD patients exhibit a defect in the import of one or more classes of peroxisomal matrix proteins into the organelle. Defects in this gene are a cause of Zellweger syndrome (ZWS), as well as peroxisome biogenesis disorder complementation group 14 (PBD-CG14), which is also known as PBD-CGJ. Alternative splicing results in multiple transcript variants.
基因ID	5824
基因名	PEX19
Swiss	P40855 (https://www.uniprot.org/uniprotkb/P40855/entry)
别名	PEX19 (YD14874),PEX19 (YD14874) Rabbit mAb,PEX19,33 kDa housekeeping protein,Peroxin-19,Peroxisomal farnesylated protein,HK33,PXF

产品验证

实验步骤

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