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FKTN (YD36108) Rabbit mAb

货号: **AYD16343**

产品信息

反应	Human, Mouse, Rat
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB IHC-P
推荐浓度	
理论分子量	54kDa
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	BxPC-3,RD,U-87MG,Mouse testis,Mouse brain,Mouse heart,Rat liver
细胞定位	Golgi apparatus membrane, Cytoplasm, Nucleus
纯化	亲和纯化

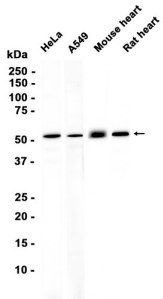
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靶点信息

研究背景	The protein encoded by this gene is a putative transmembrane protein that is localized to the cis-Golgi compartment, where it may be involved in the glycosylation of alpha-dystroglycan in skeletal muscle. The encoded protein is thought to be a glycosyltransferase and could play a role in brain development. Defects in this gene are a cause of Fukuyama-type congenital muscular dystrophy (FCMD), Walker-Warburg syndrome (WWS), limb-girdle muscular dystrophy type 2M (LGMD2M), and dilated cardiomyopathy type 1X (CMD1X). Alternatively spliced transcript variants have been found for this gene.
基因ID	2218
基因名	FKTN
Swiss	O75072 (https://www.uniprot.org/uniprotkb/O75072/entry)
别名	FKTN (YD36108),FKTN (YD36108) Rabbit mAb,FKTN,Fukutin,Fukuyama-type congenital muscular dystrophy protein,Ribitol-5-phosphate transferase,FCMD

产品验证



实验步骤

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