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ADA (YD13412) Rabbit mAb

货号: **AYD15587**

产品信息

反应	Human,Rat
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB
推荐浓度	
理论分子量	41kDa
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	HeLa,A-549
细胞定位	Cell membrane, Cell junction, Cytoplasmic vesicle lumen, Cytoplasm, Lysosome
纯化	亲和纯化

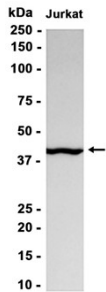
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靶点信息

研究背景	This gene encodes an enzyme that catalyzes the hydrolysis of adenosine to inosine. Various mutations have been described for this gene and have been linked to human diseases. Deficiency in this enzyme causes a form of severe combined immunodeficiency disease (SCID), in which there is dysfunction of both B and T lymphocytes with impaired cellular immunity and decreased production of immunoglobulins, whereas elevated levels of this enzyme have been associated with congenital hemolytic anemia.
基因ID	100
基因名	ADA
Swiss	P00813 (https://www.uniprot.org/uniprotkb/P00813/entry)
别名	ADA (YD13412),ADA (YD13412) Rabbit mAb,ADA,Adenosine aminohydrolase,ADA1

产品验证



实验步骤

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