

# ACAD9 Rabbit pAb

货号: **AYP13289**

## 产品信息

反应	Human,Mouse
宿主	Rabbit
克隆性	Polyclonal
预测反应	<b>WB:</b> Mus musculus
应用	<a href="#">WB</a> <a href="#">IF/ICC</a>
推荐浓度	<b>WB:</b> 1:500 - 1:2000 <b>IF/ICC:</b> 1:50 - 1:100
理论分子量	68kDa
实测分子量	69kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	SKOV3,U-251MG,MCF7,293T,NCI-H460,Mouse kidney,Mouse liver,Mouse brain
细胞定位	Mitochondrion
纯化	Affinity purification

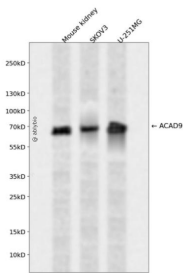
## 抗原信息

抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 1-270 of human ACAD9 (NP_054768.2).
序列	MSGCGLFLRRTTAAARACRGLVSTANRRLLRTSPPVRAFANELFLGKIKKKEVFPFPEVSQDELNEINQFLGPVEKFFTEEVD SRKIDQEGKIPDETLEKLSLGLFGLQVPEEYGGGLGFSNTMYSRLGEIISMDGSITVTLAAHQAIGLKGIIAGTTEEQKAKYLP KLASGEHIAAFCLTEPASGSDAASIRSRTLSEDKKHYYLNGSKVWITNGGLANIFTVFAKTEVVSDSDGSVKDKITAFIVERD FGGVTNGKPEDKLGIRGSNT

## 靶点信息

研究背景	This gene encodes a member of the acyl-CoA dehydrogenase family. Members of this family of proteins localize to the mitochondria and catalyze the rate-limiting step in the beta-oxidation of fatty acyl-CoA. The encoded protein is specifically active toward palmitoyl-CoA and long-chain unsaturated substrates. Mutations in this gene cause acyl-CoA dehydrogenase family member type 9 deficiency. Alternate splicing results in multiple transcript variants.
基因ID	28976
基因名	ACAD9
Swiss	Q9H845
别名	ACAD9;NPD002

## 产品验证



Western blot analysis of ACAD9 expressed in Mouse kidney, SKOV3, U-251MG using ACAD9 Rabbit pAb at 1:1000. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:5000. Lysates/proteins: 30ug per lane. Blocking buffer: 5% non-fat dry milk in TBST. Detection: ECL Enhanced Kit. Exposure time: 120s.

## 实验步骤

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