

# EHHADH Rabbit pAb

货号: B12371

## 产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Polyclonal
预测反应	<b>WB:</b> Mus musculus , Gallus gallus
应用	<a href="#">WB</a>
推荐浓度	<b>WB:</b> 1:500 - 1:2000
理论分子量	69kDa/79kDa
实测分子量	79kDa
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	SW480,A-549,HeLa,HepG2,Mouse small intestine,Rat liver,Rat kidney
细胞定位	Peroxisome
纯化	Affinity purification

## 抗原信息

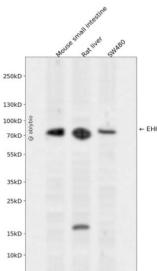
抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 444-723 of human EHHADH (NP_001957.2).
序列	EVIPSQYSSPTTIATVMNLSKKIKKIGVVVGNCFGFVGNRMLNPYYNQAYFLLEEGSKPEEV DQVLEEFGF KMGPF RVS DLA GLDVGWKS RKG QGLT GPT LPGTP ARK RGN RRYCPIPD VLCELGR FGQKTGKG WYQYDKPLGRIHKPDPWLSKFLS RYR KTHHIEPRTISQDEI LERCL YSLINEAFRILGE GIAASPEHIDVVYLHG YGWPRHKG GPMFYASTVGLPTVLEKLQKYYRQNPD IPQLEPSDYLKKLASQGNPLKEWQSLAGSPSSKL

## 靶点信息

研究背景	The protein encoded by this gene is a bifunctional enzyme and is one of the four enzymes of the peroxisomal beta-oxidation pathway. The N-terminal region of the encoded protein contains enoyl-CoA hydratase activity while the C-terminal region contains 3-hydroxyacyl-CoA dehydrogenase activity. Defects in this gene are a cause of peroxisomal disorders such as Zellweger syndrome. Two transcript variants encoding different isoforms have been found for this gene.
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基因ID	1962
基因名	EHHADH
Swiss	Q08426
别名	EHHADH;ECHD;FRTS3;L-PBE;LBFP;LBP;PBFE

## 产品验证



Western blot analysis of EHHADH expressed in Mouse small intestine, Rat liver, SW480 using EHHADH 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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