

EBP Rabbit pAb

货号: **B18771**

产品信息

反应	Mouse,Rat
以 应	Mouse, nat
宿主	Rabbit
克隆性	Polyclonal
预测反应	
应用	WB
推荐浓度	WB: 1:500 - 1:2000
理论分子量	26kDa
实测分子量	26kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.01% thiomersal,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	Mouse liver,Rat liver
细胞定位	Endoplasmic reticulum membrane,Multi-pass membrane protein
纯化	Affinity purification

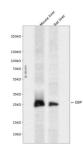
抗原信息

抗原信息	A synthetic peptide corresponding to a sequence within amino acids 50-150 of human EBP (NP_006570.1) .	
序列	SGRAAVVPLGTWRRLSLCWFAVCGFIHLVIEGWFVLYYEDLLGDQAFLSQLWKEYAKGDSRYILGDNFTVCMETITACLW GPLSLWVVIAFLRQHPLRFIL	

靶点信息

研究背景	The protein encoded by this gene is an integral membrane protein of the endoplasmic reticulum. It is a hi gh affinity binding protein for the antiischemic phenylalkylamine Ca2+ antagonist [3H]emopamil and the photoaffinity label [3H]azidopamil. It is similar to sigma receptors and may be a member of a superfamily of high affinity drug-binding proteins in the endoplasmic reticulum of different tissues. This protein shares structural features with bacterial and eukaryontic drug transporting proteins. It has four putative transme mbrane segments and contains two conserved glutamate residues which may be involved in the transport of cationic amphiphilics. Another prominent feature of this protein is its high content of aromatic amino acid residues (>23%) in its transmembrane segments. These aromatic amino acid residues have been su ggested to be involved in the drug transport by the P-glycoprotein. Mutations in this gene cause Chondro dysplasia punctata 2 (CDPX2; also known as Conradi-Hunermann syndrome).
基因 ID	10682
基因名	EBP
Swiss	Q15125
别名	EBP;CDPX2;CHO2;CPX;CPXD;MEND

产品验证



Western blot analysis of EBP expressed in Mouse liver,Rat liver using EBP Rabbit pAb at 1:1000. Secondar y 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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