

Rabbit Anti-HSD17B4 antibody

SL11296R

Product Name:	HSD17B4
Chinese Name:	羟 基 类 固醇 (17β)脱氢酶4/17β-HSD4 抗体
Alias:	hydroxysteroid (17-beta) dehydrogenase 4; 12-alpha-trihydroxy-5-beta-cholest-24-enoyl-CoA hydratase; 17 beta HSD 4; 17 beta HSD IV; 17 beta hydroxysteroid dehydrogenase 4; 17-beta-HSD 4; 17-beta-hydroxysteroid dehydrogenase 4; 17beta estradiol dehydrogenase type IV; 3 alpha 7 alpha12 alpha trihydroxy 5 beta cholest 24 enoyl CoA hydratase antibody 3-alpha; 7-alpha; Beta hydroxyacyl dehydrogenase; Beta keto reductase; D 3 hydroxyacyl CoA dehydratase; D bifunctional protein; D bifunctional protein peroxisomal; D-bifunctional protein; DBP; DHB4_HUMAN; EDH17B4; Enoyl-CoA hydratase 2; Hsd17b4; MFE 2; MFE-2; MPF-2; Multifunctional protein 2; Peroxisomal multifunctional enzyme type 2; Peroxisomal multifunctional protein 2; SDR8C1; Short chain dehydrogenase/reductase family 8C member 1; 17β-HSD4.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Chicken, Pig, Cow, Horse, Sheep, Guinea Pig, Zebrafish
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	47/80kDa
Cellular localization:	cytoplasmic Mitochondrion
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human HSD17B4 Enoyl-CoA hydratase 2:521-620/736
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized

	antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of
	antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
	17Beta-HSD4 (17Beta-hydroxysteroid dehydrogenase type 4) is also known as peroxisomal multifunctional enzyme/protein 2 (MFE-2/MFP-2), D-bifunctional enzyme or 17-Beta Estradiol dehydrogenase type IV. It belongs to the 17Beta-HSD family of proteins that regulate the availability of steroids within various tissues throughout the body. 17Beta-HSD4 inactivates Estradiol through its oxidative activity but it is primarily involved in peroxisomal fatty acid and cholesterol Beta-oxidation. It has a multi-domain structure: the dehydrogenase domain is fused to a hydratase and a lipid transfer domain. 17Beta-HSD4 is a target protein of chromeceptin and it is essential for the downstream activation of Stat6. 17Beta-HSD4-deficient patients exhibit Zellweger-like syndrome and die within the first year of life. They display neuronal migration defects, facial dysmorphisms, severe hypotonia and convulsions in the neonatal period.
	Function: Bifunctional enzyme acting on the peroxisomal beta-oxidation pathway for fatty acids. Catalyzes the formation of 3-ketoacyl-CoA intermediates from both straight-chain and 2-methyl-branched-chain fatty acids.
	Subcellular Location: Peroxisome.
Product Detail:	Tissue Specificity:
Troduct Detail.	Present in many tissues with highest concentrations in liver, heart, prostate and testis.
	DISEASE:
	Defects in HSD17B4 are a cause of D-bifunctional protein deficiency (DBPD) [MIM:261515]. DBPD is a disorder of peroxisomal fatty acid beta-oxidation.
	Similarity: Belongs to the short-chain dehydrogenases/reductases (SDR) family. Contains 1 MaoC-like domain. Contains 1 SCP2 domain.
	SWISS: P51659
	Gene ID: 3295
	Database links:

Entrez Gene: 3295Human

Entrez Gene: 15488Mouse

Entrez Gene: 79244Rat

Omim: 601860Human

SwissProt: P51659Human

SwissProt: P51660Mouse

SwissProt: P97852Rat

Unigene: 406861 Human

Unigene: 277857 Mouse

Unigene: 2082Rat

Important Note:

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.