

HEM6 rabbit pAb

Catalog No :	YT6554
Reactivity :	Human;Mouse;Rat
Applications :	WB
Target :	HEM6
Fields :	>>Porphyrin metabolism;>>Metabolic pathways;>>Biosynthesis of cofactors
Gene Name :	CPOX CPO CPX
Protein Name :	HEM6
Human Gene Id :	1371
Human Swiss Prot No :	P36551
Mouse Gene Id :	12892
Mouse Swiss Prot No :	P36552
Rat Gene Id :	304024
Rat Swiss Prot No :	Q3B7D0
Immunogen :	Synthesized peptide derived from human HEM6 AA range: 265-315
Specificity :	This antibody detects endogenous levels of HEM6 at Human/Mouse/Rat
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Polyclonal, Rabbit,IgG
Dilution :	WB 1[?]500-2000
Purification :	The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration : 1 mg/ml

Storage Stability : -15°C to -25°C/1 year (Do not lower than -25°C)

Molecularweight : 50kD

Background : The protein encoded by this gene is the sixth enzyme of the heme biosynthetic pathway. The encoded enzyme is soluble and found in the intermembrane space of mitochondria. This enzyme catalyzes the stepwise oxidative decarboxylation of coproporphyrinogen III to protoporphyrinogen IX, a precursor of heme. Defects in this gene are a cause of hereditary coproporphyria (HCP). [provided by RefSeq, Oct 2009],

Function : catalytic activity: Coproporphyrinogen-III + O(2) + 2 H(+) = protoporphyrinogen-IX + 2 CO(2) + 2 H(2)O., disease: Defects in CPOX are the cause of hereditary coproporphyria (HCP) [MIM:121300]. HCP is an acute hepatic porphyria and an autosomal dominant disease characterized by neuropsychiatric disturbances and skin photosensitivity. Biochemically, there is an overexcretion of coproporphyrin III in the urine and in the feces. HCP is clinically characterized by attacks of abdominal pain, neurological disturbances, and psychiatric symptoms. The symptoms are generally manifested with rapid onset, and can be precipitated by drugs, alcohol, caloric deprivation, infection, endocrine factors or stress. A severe variant form is harderoporphyria, which is characterized by earlier onset attacks, massive excretion of harderoporphyryn in the feces, and a marked decrease of coproporphyrinogen IX oxidase

Subcellular Location : Mitochondrion intermembrane space.

Products Images

