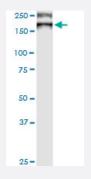


ATP7B monoclonal antibody (M01), clone 3E10

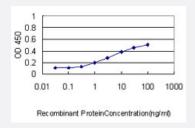
Catalog # H00000540-M01 Size 100 ug

Applications



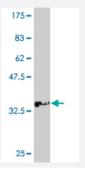
Western Blot (Tissue lysate)

ATP7B monoclonal antibody (M01), clone 3E10. Western Blot analysis of ATP7B expression in human colon.



Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged ATP7B is approximately 0.3ng/ml as a capture antibody.



Western Blot detection against Immunogen (36.08 KDa).

Specification

Product Description

Mouse monoclonal antibody raised against a partial recombinant ATP7B.



Product Information

Immunogen	ATP7B (NP_000044, 1372 a.a. ~ 1465 a.a) partial recombinant protein with GST tag. MW of the GS T tag alone is 26 KDa.
Sequence	QLKCYKKPDLERYEAQAHGHMKPLTASQVSVHIGMDDRWRDSPRATPWDQVSYVSQVSLSSLT SDKPSRHSAAADDDGDKWSLLLNGRDEEQYI
Host	Mouse
Reactivity	Human
Interspecies Antigen Sequence	Mouse (85); Rat (84)
Isotype	lgG1 Kappa
Quality Control Testing	Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (36.08 KDa).
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

Western Blot (Tissue lysate)

ATP7B monoclonal antibody (M01), clone 3E10. Western Blot analysis of ATP7B expression in human colon.

Protocol Download

Western Blot (Recombinant protein)

Protocol Download

Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged ATP7B is approximately 0.3ng/ml as a capture antibody.

Protocol Download

ELISA

Gene Info — ATP7B

Entrez GenelD <u>540</u>



Product Information

GeneBank Accession#	<u>NM_000053</u>
Protein Accession#	NP_000044
Gene Name	ATP7B
Gene Alias	PWD, WC1, WD, WND
Gene Description	ATPase, Cu++ transporting, beta polypeptide
Omim ID	<u>277900</u> <u>606882</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene is a member of the P-type cation transport ATPase family and encodes a protein with s everal membrane-spanning domains, an ATPase consensus sequence, a hinge domain, a phosp horylation site, and at least 2 putative copper-binding sites. This protein functions as a monomer, exporting copper out of the cells, such as the efflux of hepatic copper into the bile. Alternate transc riptional splice variants, encoding different isoforms with distinct cellular localizations, have been characterized. Mutations in this gene have been associated with Wilson disease (WD). [provided by RefSeq
Other Designations	ATPase, Cu(2+)- transporting, beta polypeptide OTTHUMP00000040880 Wilson disease-associ

Publication Reference

Characterization of Sandwich-Cultured Hepatocytes as an In Vitro Model to Assess the Hepatobiliary
 Disposition of Copper.

Ansede JH, Wright MR, St Claire RL, Hart RW, Gefroh HA, Brouwer KR.

Drug Metabolism and Disposition 2009 May; 37(5):969.

Application: WB, Rat, dog, human, Hepatocytes

Disease

- Chromosome Aberrations
- Genetic Predisposition to Disease
- Hepatolenticular Degeneration
- Kidney Failure
- Liver Failure





- Mental Disorders
- Motor Skills