Leader in Biomolecular Solutions for Life Science

CYP51A1 Rabbit mAb

Catalog No.: A23370 Recombinant 1 Publications



Basic Information

Observed MW 55kDa

Calculated MW 57kDa

Category Primary antibody

Applications ELISA,WB,IHC-P

Cross-Reactivity Human, Mouse, Rat

CloneNo number ARC59516

Recommended Dilutions

Background

This gene encodes a member of the cytochrome P450 superfamily of enzymes. The cytochrome P450 proteins are monooxygenases which catalyze many reactions involved in drug metabolism and synthesis of cholesterol, steroids and other lipids. This endoplasmic reticulum protein participates in the synthesis of cholesterol by catalyzing the removal of the 14alpha-methyl group from lanosterol. Homologous genes are found in all three eukaryotic phyla, fungi, plants, and animals, suggesting that this is one of the oldest cytochrome P450 genes. Two transcript variants encoding different isoforms have been found for this gene.

Immunogen Information

WB	1:1000 - 1:5000	Gene ID	Swiss Prot	
IHC-P	1:50 - 1:200	1595	Q16850	

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 400-500 of human CYP51A1 (NP_000777.1).

Synonyms

LDM; CP51; CYP51; CYPL1; P450L1; P450-14DM; CYP51A1

Contact

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Product Information

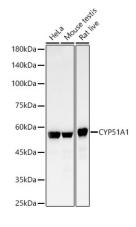
Source Rabbit

Isotype lgG

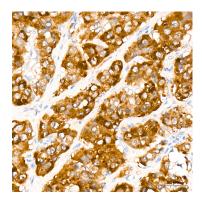
Purification Affinity purification

Storage

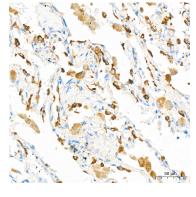
Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.



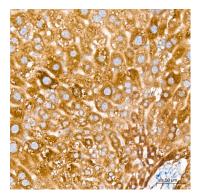
Western blot analysis of various lysates, using CYP51A1 Rabbit mAb (A23370) at 1:2000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 30s.



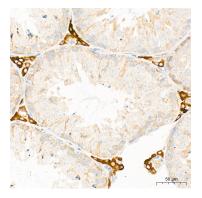
Immunohistochemistry analysis of paraffinembedded Human liver cancer tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



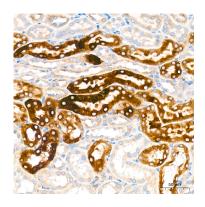
Immunohistochemistry analysis of paraffinembedded Human lung tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



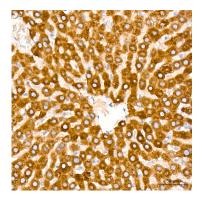
Immunohistochemistry analysis of paraffinembedded Mouse liver tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse testis tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.

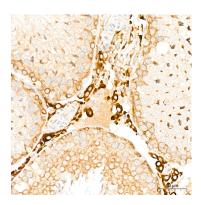


Immunohistochemistry analysis of paraffinembedded Mouse kidney tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat liver tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.

Validation Data



Immunohistochemistry analysis of paraffinembedded Rat testis tissue using CYP51A1 Rabbit mAb (A23370) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.