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Datasheet for ABIN7300598 anti-CYP2S1 antibody (Center)

3 Images



Overview

Quantity:	100 µL
Target:	CYP2S1
Binding Specificity:	Center
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CYP2S1 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF), Immunochromatography (IC)

Product Details

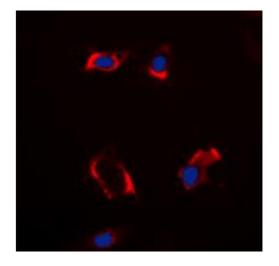
Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human Cytochrome P450 2S1.
Specificity:	Recognizes endogenous levels of Cytochrome P450 2S1 protein.
Characteristics:	Rabbit polyclonal antibody to Cytochrome P450 2S1
Purification:	The antibody was purified by immunogen affinity chromatography.
Target Details	

Target:	CYP2S1
Alternative Name:	Cytochrome P450 2S1 (CYP2S1 Products)

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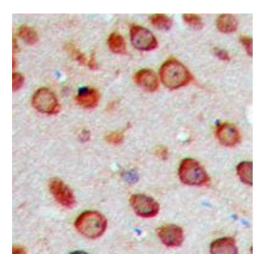
Target Details	
Background:	Cytochrome P450 2S1, CYPIIS1
Gene ID:	29785
UniProt:	Q96SQ9
Application Details	
Application Notes:	WB (1:500 - 1:1000), IH (1:100 - 1:200), IF/IC (1:100 - 1:500)
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.
Buffer: Preservative:	
	0.01 % sodium azide.
Preservative:	0.01 % sodium azide. Sodium azide
Preservative:	0.01 % sodium azide. Sodium azide This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which
Preservative: Precaution of Use:	0.01 % sodium azide. Sodium azide This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

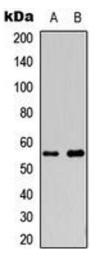
Images



Immunofluorescence

Image 1. Immunofluorescent analysis of Cytochrome P450 2S1 staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI





was used to stain the cell nuclei (blue).

Immunohistochemistry

Image 2. Immunohistochemical analysis of Cytochrome P450 2S1 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

Western Blotting

Image 3. Western blot analysis of Cytochrome P450 2S1 expression in HepG2 (A), COLO205 (B) whole cell lysates.

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