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## anti-CYP1B1 antibody (Internal Region)

3 Images



Go to Product page

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Quantity:	100 μL
Target:	CYP1B1
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CYP1B1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human CYP1B1, corresponding to a region within the internal amino acids.

Immunogen:	A synthesized peptide derived from human CYP1B1, corresponding to a region within the internal amino acids.	
Isotype:	IgG	
Specificity:	CYP1B1 Antibody detects endogenous levels of total CYP1B1.	
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog,Chicken	
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).	

#### **Target Details**

Target:	CYP1B1	

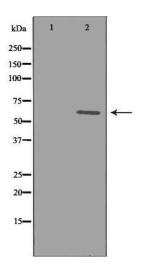
### **Target Details**

Alternative Name:	CYP1B1 (CYP1B1 Products)		
Background:	Description: Cytochromes P450 are a group of heme-thiolate monooxygenases. In liver		
	microsomes, this enzyme is involved in an NADPH-dependent electron transport pathway. It		
	oxidizes a variety of structurally unrelated compounds, including steroids, fatty acids, retinoid		
	and xenobiotics. Preferentially oxidizes 17beta-estradiol to the carcinogenic 4-hydroxy		
	derivative, and a variety of procarcinogenic compounds to their activated forms, including		
	polycyclic aromatic hydrocarbons. Promotes angiogenesis by removing cellular oxygenation		
	products, thereby decreasing oxidative stress, release of antiangiogenic factor THBS2, then		
	allowing endothelial cells migration, cell adhesion and capillary morphogenesis. These changes		
	are concommitant with the endothelial nitric oxide synthase activity and nitric oxide synthesis.		
	Plays an important role in the regulation of perivascular cell proliferation, migration, and surviva		
	through modulation of the intracellular oxidative state and NF-kappa-B expression and/or		
	activity, during angiogenesis. Contributes to oxidative homeostasis and ultrastructural		
	organization and function of trabecular meshwork tissue through modulation of POSTN		
	expression.		
	Gene: CYP1B1		
Molecular Weight:	61kDa		
Gene ID:	1545		
UniProt:	Q16678		
Pathways:	Steroid Hormone Biosynthesis		
Application Details			
Application Notes:	WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000		
Restrictions:	For Research Use only		
Handling			
Format:	Liquid		
Concentration:	1 mg/mL		
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 %		
	glycerol.		
Preservative:	Sodium azide		

#### Handling

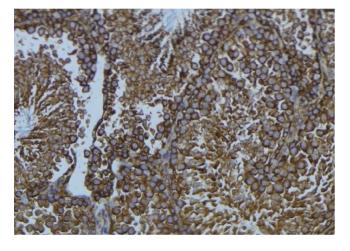
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months

#### **Images**



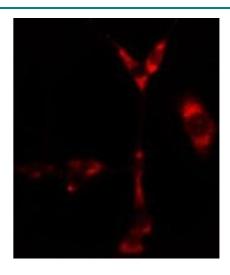
#### **Western Blotting**

**Image 1.** Western blot analysis of extracts of CEM, using CYP1B1 antibody. The lane on the left is treated with the antigen-specific peptide.



#### **Immunohistochemistry**

**Image 2.** ABIN6276665 at 1/100 staining Mouse testis tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22¡ãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary



#### Immunofluorescence (fixed cells)

**Image 3.** ABIN6276665 staining Hela cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25;ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37;ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod