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anti-CYP26A1 antibody (Internal Region)

2 Images



Overview	
Quantity:	100 μL
Target:	CYP26A1
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CYP26A1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunocytochemistry (ICC), Immunofluorescence (IF)
Product Details	
Immunogen:	A synthesized peptide derived from human CYP26A1, corresponding to a region within the internal amino acids.
Isotype:	lgG
Specificity:	CYP26A1 Antibody detects endogenous levels of total CYP26A1.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	CYP26A1

Target Details

Target Details	
Alternative Name:	CYP26A1 (CYP26A1 Products)
Background:	Description: Plays a key role in retinoic acid metabolism. Acts on retinoids, including all-trans- retinoic acid (RA) and its stereoisomer 9-cis-RA. Capable of both 4-hydroxylation and 18- hydroxylation. Responsible for generation of several hydroxylated forms of RA, including 4-OH- RA, 4-oxo-RA and 18-OH-RA. Gene: CYP26A1
Molecular Weight:	56 kDa
Gene ID:	1592
UniProt:	043174
Pathways:	Retinoic Acid Receptor Signaling Pathway, Monocarboxylic Acid Catabolic Process
Application Details	
Application Notes:	WB 1:500-1:1000, 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
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Store at -20 °C. Stable for 12 months from date of receipt.

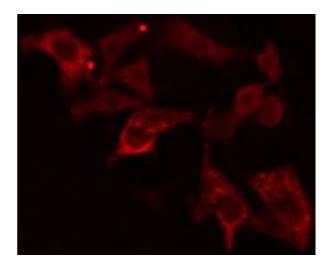
-20 °C

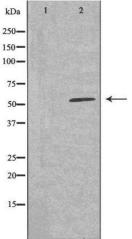
12 months

Storage:

Expiry Date:

Storage Comment:





Immunofluorescence (fixed cells)

Image 1. ABIN6274640 staining HepG2 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°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibody.

Western Blotting

Image 2. Western blot analysis of extracts from HepG2 cells using Cytochrome P450 26A1 antibody. The lane on the left is treated with the antigen-specific peptide.