antibodies -online.com







anti-QPRT antibody (AA 113-254)

Images



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Quantity:	100 μg
Target:	QPRT
Binding Specificity:	AA 113-254
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This QPRT antibody is un-conjugated
Application:	Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF)
Product Details	
Immunogen:	Recombinant Human Nicotinate-nucleotide pyrophosphorylase [carboxylating] protein (113-254AA)
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified
Target Details	
Target:	QPRT
Alternative Name:	QPRT (QPRT Products)
Background:	Background: Involved in the catabolism of quinolinic acid (QA).

Target Details

Aliases: NADC_HUMAN antibody, Nicotinate nucleotide pyrophosphorylase (carboxylating) antibody, Nicotinate-nucleotide pyrophosphorylase [carboxylating] antibody, QAPRTase antibody, QPRT antibody, QPRTase antibody, Quinolinate phosphoribosyltransferase [decarboxylating] antibody, Quinolinate phosphoribosyltransferase antibody

UniProt:

Q15274

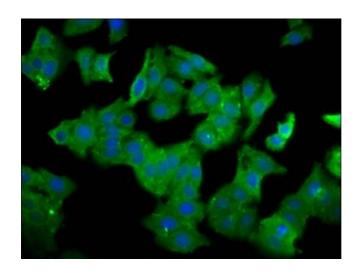
Application Details

Application Notes:	Recommended dilution: IHC:1:200-1:500, IF:1:50-1:200,
Restrictions:	For Research Use only

Handling

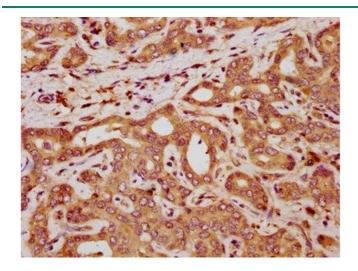
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

Images



Immunofluorescence

Image 1. Immunofluorescence staining of HepG2 cells with ABIN7161592 at 1:100, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Immunohistochemistry

Image 2. IHC image of ABIN7161592 diluted at 1:300 and staining in paraffin-embedded human liver cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.