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Datasheet for ABIN7298622 anti-HKR1 antibody (Center)

3 Images



Overview

Quantity:	100 µL
Target:	HKR1
Binding Specificity:	Center
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This HKR1 antibody is un-conjugated
Application:	Western Blotting (WB), Immunofluorescence (IF), Immunohistochemistry (IHC), Immunoprecipitation (IP), Immunochromatography (IC)

Product Details

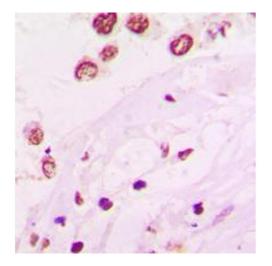
Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human HKR1.
Specificity:	Recognizes endogenous levels of HKR1 protein.
Characteristics:	Rabbit polyclonal antibody to HKR1
Purification:	The antibody was purified by immunogen affinity chromatography.
Target Details	
Targati	

Target:	HKR1
Alternative Name:	HKR1 (HKR1 Products)

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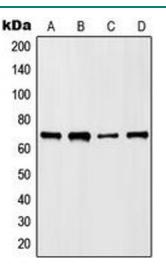
Target Details	
Background:	ZNF875, Krueppel-related zinc finger protein 1, Protein HKR1, Zinc finger protein 875
Gene ID:	284459
UniProt:	P10072
Application Details	
Application Notes:	WB (1:500 - 1:1000), IH (1:100 - 1:200), IF/IC (1:100 - 1:500), IP (1:10 - 1:100)
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.
Buffer: Preservative:	
	0.01 % 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.

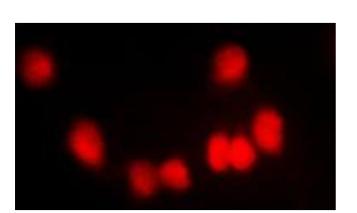
Images



Immunohistochemistry

Image 1. Immunohistochemical analysis of HKR1 staining in human lung cancer 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 conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.





Western Blotting

Image 2. Western blot analysis of HKR1 expression in HepG2 (A), NIH3T3 (B), mouse liver (C), rat liver (D) whole cell lysates.

Immunofluorescence

Image 3. Immunofluorescent analysis of HKR1 staining in NIH3T3 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 humidified 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).

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