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Datasheet for ABIN7296780 anti-RAD23B antibody (N-Term)

3 Images



Overview

Quantity:	100 µL
Target:	RAD23B
Binding Specificity:	N-Term
Reactivity:	Human, Mouse, Rat, Cow, Monkey
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This RAD23B antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF), Immunoprecipitation (IP), Immunochromatography (IC)

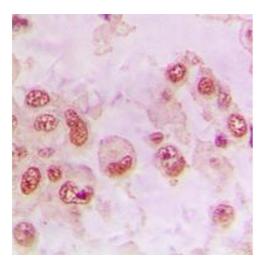
Product Details

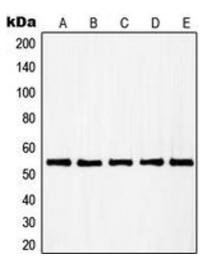
Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the N-term region of human RAD23B.
Specificity:	Recognizes endogenous levels of RAD23B protein.
Characteristics:	Rabbit polyclonal antibody to RAD23B
Purification:	The antibody was purified by immunogen affinity chromatography.
Target Details	

Target:	RAD23B
Alternative Name:	RAD23B (RAD23B Products)

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Target Details		
Background:	UV excision repair protein RAD23 homolog B, HR23B, hHR23B, XP-C repair-complementing complex 58 kDa protein, p58	
Gene ID:	5887, 19359, 298012	
UniProt:	P54727, P54728, Q4KMA2	
Pathways:	DNA Damage Repair	
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.	
Preservative:	Sodium azide	
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:	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.	
Expiry Date:	12 months	



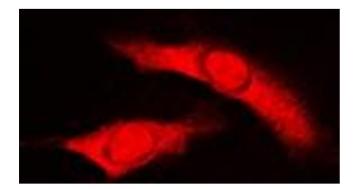


Immunohistochemistry

Image 1. Immunohistochemical analysis of RAD23B 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 RAD23B expression in A431 (A), HuvEc (B), HEK293 (C), HeLa (D), HT29 (E) whole cell lysates.



Immunofluorescence

Image 3. Immunofluorescent analysis of RAD23B staining in HuvEc 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 594conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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