

Product Data Sheet

Anti-RAD23B Antibody

Catalog #	Source	Reactivity	Applications		
CPA1982	Rabbit	H, M, R, B, Mk	WB, IH, IF/IC		
Description	Rab	bit polyclonal antibody to	o RAD23B		
Immunogen	KLH	-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	regi	on of human RAD23B. Th	ne exact sequence is proprietary.		
Purification	The	antibody was purified by	/ immunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous leve	ls of RAD23B protein.		
Clonality	Poly	rclonal			
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	RAD	23B			
Alternative Na	ames UV	excision repair protein R/	AD23 homolog B; HR23B; hHR23B; XP-C		
	repa	air-complementing comp	lex 58 kDa protein; p58		
Entrez Gene	588	7 (Human); 19359 (Mous	se); 298012 (Rat)		
SwissProt	P54	727 (Human); P54728 (N	louse); Q4KMA2 (Rat)		
Storage/Stabi	lity Ship	ped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	free	ze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

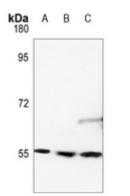
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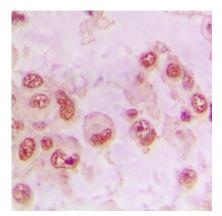


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Western blot analysis of RAD23B expression in HEK293T (A), K562 (B), A549 (C) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 55 kD)



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.



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 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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