

Product Data Sheet

Anti-XPA Antibody

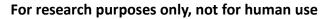
Catalog #	Source	Reactivity	Applications		
CPA2233	Rabbit	H, M, R, D	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody	to XPA		
Immunogen		KLH-conjugated synthetic pe	eptide encompassing a sequence within the C-term		
	region of human XPA. The exact sequence is proprietary.				
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity	Specificity Recognizes endogenous levels of XPA protein.				
Clonality		Polyclonal			
Conjugation					
Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glyce					
		and 0.01% sodium azide.			
Dilution WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 -) - 1/100), IF/IC (1/50 - 1/200)			
Gene Symbol		ХРА			
Alternative Na	ames	XPAC; DNA repair protein co	mplementing XP-A cells; Xeroderma pigmentosum		
		group A-complementing pro	otein		
Entrez Gene		7507 (Human); 22590 (Mouse)			
SwissProt		P23025 (Human); Q64267 (I	Mouse)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
		freeze/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

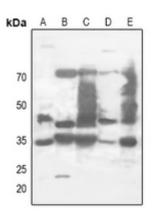
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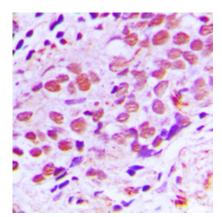




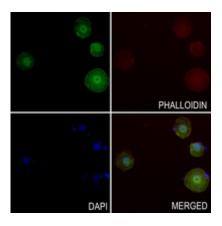
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Western blot analysis of XPA expression in HepG2 (A), mouse liver (B), mouse spleen (C), rat liver (D), rat spleen (E) whole cell lysates. (Predicted band size: 31 kD; Observed band size: 40 kD)



Immunohistochemical analysis of XPA staining in human breast 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 XPA staining in Hela 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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