

Product Data Sheet

Anti-Topoisomerase 2 beta Antibody

Catalog #	Source	Reactivity	Applications		
CPA2183	Rabbit	H, M, D, Mk, P	WB, IH, ChIP		
Description	Rabl	Rabbit polyclonal antibody to Topoisomerase 2 beta			
Immunogen	KLH	conjugated synthetic peptide encom	passing a sequence within the N-term		
	regio	on of human Topoisomerase 2 beta.	The exact sequence is proprietary.		
Purification	The	antibody was purified by immunoge	n affinity chromatography.		
Specificity	Reco	ognizes endogenous levels of Topoisc	merase 2 beta protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium phosphate, 0.8	87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/50 - 1/100), ChI	P (1/10 - 1/100)		
Gene Symbol	ТОР	2B			
Alternative Na	ames DNA	topoisomerase 2-beta; DNA topoiso	merase II, beta isozyme		
Entrez Gene	715	5 (Human)			
SwissProt	Q02	880 (Human)			
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery aliquot and	d 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

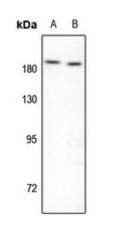
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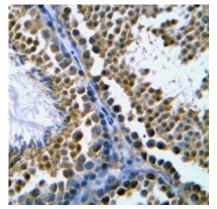
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Western blot analysis of Topoisomerase 2 beta expression in HEK293T (A), mouse testis (B) whole cell lysates. (Predicted band size: 183 kD; Observed band size: 183 kD)



Immunohistochemical analysis of Topoisomerase 2 beta staining in human tonsil 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.

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