

## **Product Data Sheet**

## Anti-DDX3X (Phospho-T322) Antibody

Catalog #	Source	Reactivity	Applications		
CPA5306	Rabbit	Н	WB, IH, IF/IC		
Description	tion Rabbit polyclonal antibody to DDX3X (Phospho-T322)				
Immunogen		KLH-conjugated synthetic p	hosphopeptide corresponding to residues surrounding		
		T322 of human DDX3X prot	ein. The exact sequence is proprietary.		
Purification		The antibody was purified I	by immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	els of DDX3X protein only when phosphorylated at T322.		
Clonality	onality Polyclonal				
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/2000), IH (1/50 - 1/200), IF/IC (1/50 - 1/100)			
Gene Symbol		DDX3X			
Alternative Names		DBX; DDX3; ATP-dependent RNA helicase DDX3X; DEAD box protein 3,			
		X-chromosomal; DEAD box,	X isoform; Helicase-like protein 2; HLP2		
Entrez Gene		1654 (Human)			
SwissProt		000571 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon delivery 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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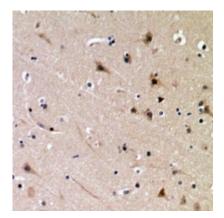
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Western blot analysis of DDX3X (Phospho-T322) expression in HeLa TNFa-treated (A) whole cell lysates. (Predicted band size: 73 kD; Observed band size: 78 kD)



Immunohistochemical analysis of DDX3X (Phospho-T322) staining in human brain 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 DDX3X (Phospho-T322) 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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