

Product Data Sheet

Anti-DDX24 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3893	Rabbit	H, M, R, Mk	WB, IH		
Description	Rabb	Rabbit polyclonal antibody to DDX24			
Immunogen	KLH-	conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	regio	region of human DDX24. The exact sequence is proprietary.			
Purification	The a	antibody was purified by	<i>immunogen affinity chromatography.</i>		
Specificity	Reco	gnizes endogenous leve	ls of DDX24 protein.		
Clonality	Polyc	clonal			
Conjugation					
Form	Liqui	d in 0.42% Potassium pl	nosphate, 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)		
Gene Symbol	DDX	24			
Alternative Na	ames ATP-0	dependent RNA helicase	e DDX24; DEAD box protein 24		
Entrez Gene	5706	2 (Human); 27225 (Mou	ise)		
SwissProt	Q9G	ZR7 (Human); Q9ESV0 (I	Mouse)		
Storage/Stabi	lity Ship	oed at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	freez	e/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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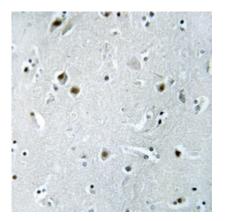
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For research purposes only, not for human use

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Western blot analysis of DDX24 expression in Hela (A), HEK293T (B) whole cell lysates. (Predicted band size: 96 kD; Observed band size: 120 kD)



Immunohistochemical analysis of DDX24 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.

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