

Product Data Sheet

Anti-DLX5 Antibody

Catalog #	Source	Reactivity	Applications
CPA1346	Rabbit	H, M, R, C, Z	WB, IH, IF/IC
Description	R	abbit polyclonal antibody to	DLX5
Immunogen	К	LH-conjugated synthetic pe	otide encompassing a sequence within the center
	re	egion of human DLX5. The e	xact sequence is proprietary.
Purification	Т	he antibody was purified by	immunogen affinity chromatography.
Specificity	R	ecognizes endogenous leve	ls of DLX5 protein.
Clonality	Р	Polyclonal	
Conjugation			
Form	Li	iquid in 0.42% Potassium pł	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	а	nd 0.01% sodium azide.	
Dilution	V	VB (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)
Gene Symbol	D	DLX5	
Alternative Na	ames H	lomeobox protein DLX-5	
Entrez Gene	1	.749 (Human); 13395 (Mous	e); 25431 (Rat)
SwissProt	Р	256178 (Human); P70396 (M	ouse); P50575 (Rat)
Storage/Stabi	lity S	hipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid
	fr	reeze/thaw cycles.	

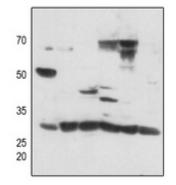
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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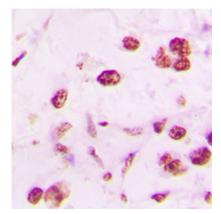
kDa A B C D E F



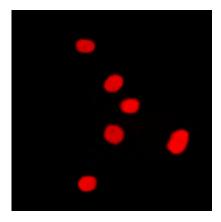
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Western blot analysis of DLX5 expression in Hela (A), A2780 (B), mouse lung (C), mouse brain (D), mouse kidney (E), rat kidney (F) whole cell lysates. (Predicted band size: 31 kD; Observed band size: 31 kD)



Immunohistochemical analysis of DLX5 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 DLX5 staining in Raw264.7 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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