

Product Data Sheet

Anti-HOXB2 Antibody

Catalog #	Source	Reactivity	Applications	
CPA1541	Rabbit	н, м, R	WB, IH	
Description		Rabbit polyclonal antibody to	O HOXB2	
Immunogen	I	KLH-conjugated synthetic pe	otide encompassing a sequence within the center	
	I	region of human HOXB2. The	exact sequence is proprietary.	
Purification	-	The antibody was purified by	immunogen affinity chromatography.	
Specificity	ļ	Recognizes endogenous leve	ls of HOXB2 protein.	
Clonality	I	Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	i	and 0.01% sodium azide.		
Dilution	,	WB (1/500 - 1/1000), IH (1/50	- 1/100)	
Gene Symbol	I	HOXB2		
Alternative Na	ames	HOX2H; Homeobox protein H	lox-B2; Homeobox protein Hox-2.8; Homeobox protein	
	I	Нох-2Н; К8		
Entrez Gene	:	3212 (Human); 103889 (Mou	ise)	
SwissProt	I	P14652 (Human); P0C1T1 (N	ouse)	
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
	1	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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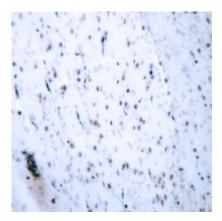
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Western blot analysis of HOXB2 expression in H446 (A), H1792 (B), mouse kidney (C), rat kidney (D), rat muscle (E) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 38 kD)



Immunohistochemical analysis of HOXB2 staining in rat 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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