

# **Product Data Sheet**

### **Anti-SOX8** Antibody

Catalog #	Source	Poostivity	Applications			
Catalog #	Source	Reactivity	Applications			
CPA6120	Rabbit	H, M, R, C, P, Z	WB, IH			
Description	Ra	bbit polyclonal antibody to	SOX8			
Immunogen	KL	H-conjugated synthetic per	otide encompassing a sequence within the center			
	re	gion of human SOX8. The e	xact sequence is proprietary.			
Purification	Th	e antibody was purified by	immunogen affinity chromatography.			
Specificity	Re	cognizes endogenous level	s of SOX8 protein.			
Clonality	Ро	lyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	an	d 0.01% sodium azide.				
Dilution	W	B (1/500 - 1/1000), IH (1/50 -	- 1/200)			
Gene Symbol	SC	0X8; SOX9; SOX17; SOX18				
Alternative Na	ames SC	X8; Transcription factor SC	X-8; SOX9; Transcription factor SOX-9; SOX17;			
	Tra	anscription factor SOX-17; S	SOX18; Transcription factor SOX-18			
Entrez Gene		30812, 6662, 64321, 54345 (Human); 20681, 20682, 20671, 20672 (Mouse)				
SwissProt	P5	7073, P48436, Q9H6I2, P3	5713 (Human); Q04886, Q04887, Q61473, P43680			
	(№	louse)				
Storage/Stabi	lity Sh	ipped at 4°C. Upon delivery	y aliquot and store at -20°C for one year. Avoid			
	fre	eeze/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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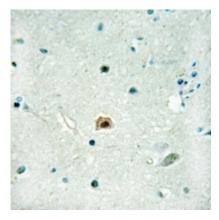
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Western blot analysis of SOX8 expression in HEK293T (A), HCT116 (B), HepG2 (C), PC12 (D), CT26 (E) whole cell lysates. (Predicted band size: 47; 56; 44; 40 kD; Observed band size: 47 kD)



Immunohistochemical analysis of SOX8 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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