

## **Product Data Sheet**

## Anti-SOX9 (Phospho-S181) Antibody

Catalog #	Source	Reactivity	Applications
CPA2094	Rabbit	H, M, R, D, P	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody to	SOX9 (Phospho-S181)
Immunogen	KLH-	conjugated synthetic pho	osphopeptide corresponding to residues surrounding
	S181	of human SOX9 protein.	The exact sequence is proprietary.
Purification	The a	intibody was purified by	immunogen affinity chromatography.
Specificity	Reco	gnizes endogenous level	s of SOX9 protein only when phosphorylated at S181.
Clonality	Polyc	lonal	
Conjugation			
Form	Liqui	d in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and (	).01% sodium azide.	
Dilution	WB (	1/500 - 1/1000), IH (1/50 -	- 1/100), IF/IC (1/50 - 1/200)
Gene Symbol	SOX9	1	
Alternative Na	ames Trans	cription factor SOX-9	
Entrez Gene	6662	(Human); 20682 (Mouse	e)
SwissProt	P484	36 (Human); Q04887 (M	ouse)
Storage/Stabi	lity Shipp	ed at 4°C. Upon delivery	v 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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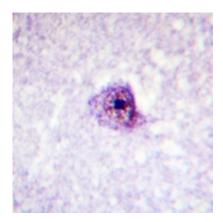
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Western blot analysis of SOX9 (Phospho-S181) expression in DLD (A), HEK293T (B), A375 (C) whole cell lysates. (Predicted band size: 56 kD; Observed band size: 56 kD)



Immunohistochemical analysis of SOX9 (Phospho-S181) 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 SOX9 (Phospho-S181) staining in A549 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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