

Product Data Sheet

Anti-SOX2 Antibody

Catalog #	Source	Reactivity	Applications		
CQA2565	Rabbit	H <i>,</i> M, R	WB, IH, IF/IC		
Description	F	Rabbit polyclonal antibody	to SOX2		
Immunogen	ŀ	KLH-conjugated synthetic p	eptide of human SOX2		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity	F	Recognizes endogenous lev	els of SOX2 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form Liquid in 0.42% Potassium phos		ا Liquid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ä	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	S	SOX2			
Alternative Na	ames	Transcription factor SOX-2			
Entrez Gene	6	6657 (Human); 20674 (Mou	ise)		
SwissProt	F	P48431 (Human); P48432 (Mouse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	f	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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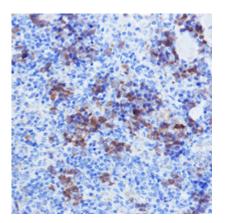
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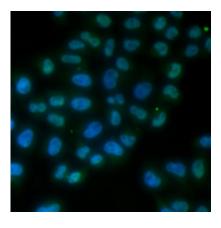
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Western blot analysis of SOX2 expression in mouse brain (A), rat brain (B) whole cell lysates. (Predicted band size: 34 kD; Observed band size: 35 kD)



Immunohistochemical analysis of SOX2 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 SOX2 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 humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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