

Anti-CREB (Phospho-S133) Antibody

Catalog #	Source	Reactivity	Applications
CQA8086	Rabbit	H, R	WB, IH, IP
Description	Rabbit monoclonal antibody to CREB (Phospho-S133)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S133 of human CREB protein. The exact sequence is proprietary.		
Purification	The antibody was purified by immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of CREB protein only when phosphorylated at S133.		
Clonality	Monoclonal		
Conjugation			
Form	Liquid in 50mM Tris-Glycine (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide and 0.05% BSA.		
Dilution	WB (1/500 - 1/1000), IH (1/50 - 1/100), IP (1/10 - 1/50)		
Gene Symbol	CREB1		
Alternative Names	Cyclic AMP-responsive element-binding protein 1; CREB-1; cAMP-responsive element-binding protein 1		
Entrez Gene	1385 (Human); 81646 (Rat)		
SwissProt	P16220 (Human); P15337 (Rat)		
Storage/Stability	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid 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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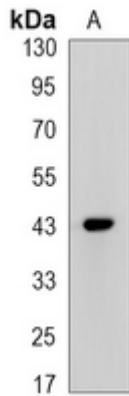
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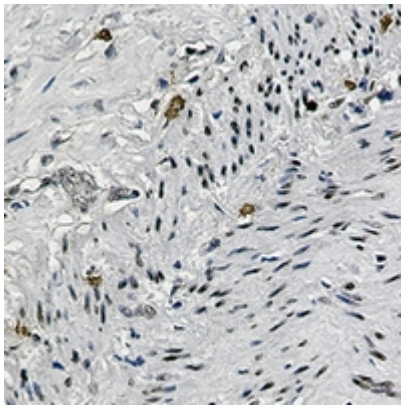
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Product Data Sheet



Western blot analysis of CREB (Phospho-S133) expression in C6 UV treated (A) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 43 kD)



Immunohistochemical analysis of CREB (Phospho-S133) 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.

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