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Application	s Detection	Clonality	Isotype
Flow Cytomet	ry N/A	Monoclonal	Rabbit IgGk
Format:	FITC		
Cross Reactivity:	Predicted to work with mouse, rat	and other homologues.	
Formulation:	1X PBS, 0.09% NaN3, 0.2% BSA		
Preparation:	Protein A+G		
Reactivity:	Human,Mouse		
Recommended Usage:	For flow cytometric staining, the suggested use of this reagent is 5 μ L per million cells or 5 μ L per 100 μ L of staining volume. It is recommended that the reagent be titrated for optimal performance for each application.		
Immunogen:	A synthetic phospho-peptide corresponding to residues surrounding Ser133 of human phospho CREB		
Description:	The cyclic AMP-responsive element (CRE)-binding protein (CREB) is a 43-kDa leucine zipper transcription factor that belongs to the CREB/ATF family and modulates the activity of many eukaryotic transcription factors by altering their subcellular localization, DNA binding, or transactivation response. It regulate downstream targets involving in various cellular functions including cell proliferation, survival, and differentiation (1-2). CREB activity is considered critical for neuronal function and regulates synaptic plasticity, long-term memory formation and neuronal survival (3). CREB serine 133 (Ser-133) residue is phosphorylated by various kinases and this phosphorylation promotes the interaction of CREB with a number of transcription coactivators, especially the histone acetyltransferases CREB-binding protein (CBP) or p300 (4,5). CREB can be phosphorylated and thus activated in response to various stimuli such as growth factors, neurotransmitters, stress signals that increase intracellular cAMP or calcium levels. CREB is also activated by phosphorylation at Ser-133 through nuclear translocation of transducer of regulated CREB activity (TORC) coactivators (6,7). CREB is phosphorylated in response to hormonal stimuli that increase intracellular cAMP production, but it can also be phosphorylated in response to a wide variety of extra-cellular signals, including growth factors, osmotic stress and ultraviolet irradiation. Several kinases including ribosomal protein S6 kinase (pp90RSK), protein kinase (CPKC), protein kinase B/AKT, and mitogen- and stress-activated protein kinase (MSK-1) can phosphorylate CREB at Ser-133 (1,8). Different growth factors such as mast/stem cell growth factor, basic fibroblast growth factor, and Granulocyte-macrophage colony-stimulating factor (GM-CSF), can all induce phosphorylation of CREB (9, 10). Phospho CREB, upon activation, dimerizes and binds to the promoter regions of its target gene that contains cAMP response element (CRE site), TGACGTCA, or CRE half sites CGTCA/TGACG, and p		



References:

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Phospho-CREB (S133) FITC

Flow cytometric analysis of SK.N.MC cells secondary antibody only negative control (blue) or untreated (red) or treated with Forskolin (green) using Phospho-CREB (Ser133) FITC conjugated antibody CREBS133-4D11. Cat. #2128.

