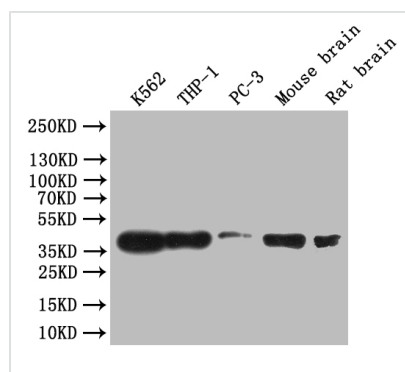




# CREB1 Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA081853A0HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P16220
<b>Immunogen</b>	A synthesized peptide derived from Human CREB1
<b>Species Reactivity</b>	Human, Mouse, Rat
<b>Tested Applications</b>	ELISA, WB, FC; Recommended dilution: WB:1:500-1:2000, FC:1:50-1:200
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	Rabbit IgG
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Recombinant Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Epigenetics and Nuclear Signaling;Immunology
<b>Gene Names</b>	CREB1
<b>Clone No.</b>	21E3

## Image

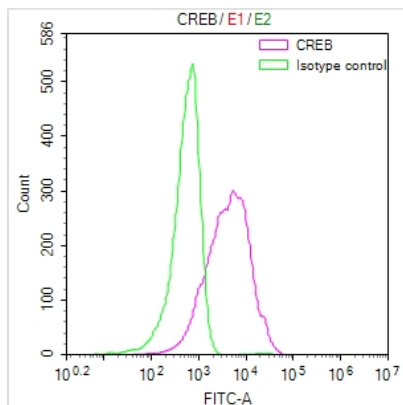


### Western Blot

Positive WB detected in: K562 whole cell lysate, THP-1 whole cell lysate, Mouse brain tissue lysate, Rat brain tissue lysate, All lanes: CREB antibody at 1:1000

### Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution  
Predicted band size: 36 kDa  
Observed band size: 36 kDa



Overlay Peak curve showing Jurkat cells stained with CSB-RA081853A0HU (red line) at 1:50. The cells were fixed in 4% formaldehyde and permeated by 0.2% TritonX-100. Then 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (1 $\mu$ g/1 $\times$ 10<sup>6</sup> cells) for 45min at 4 $^{\circ}$ C. The secondary antibody used was FITC-conjugated Goat Anti-rabbit IgG(H+L) at 1:200 dilution for 35min at 4 $^{\circ}$ C. Control antibody (green line) was rabbit IgG (1 $\mu$ g/1 $\times$ 10<sup>6</sup> cells) used under the same conditions. Acquisition of >10,000 events was performed.

## Description

CUSABIO's approach to developing a recombinant monoclonal antibody against CREB1 commenced with the immunization of a rabbit using a synthesized peptide from human CREB1 protein. B cells were subsequently isolated from the immunized rabbit, and RNA was extracted from these B cells. The extracted RNA was reverse-transcribed into cDNA, which served as a template for extending CREB1 antibody genes using degenerate primers. The engineered CREB1 antibody genes were incorporated into a plasmid vector and transfected into host cells for expression. The resulting CREB1 recombinant monoclonal antibody was then purified from the cell culture supernatant using affinity chromatography. Its suitability for ELISA, WB, and FC applications was confirmed, demonstrating specific reactivity with CREB1 proteins from human, mouse, and rat species.

CREB1 is a crucial transcription factor that regulates gene expression in response to a wide range of signals and stimuli. Its roles extend to various physiological processes, including learning and memory, cell growth and differentiation, metabolism, and the cellular response to stress.