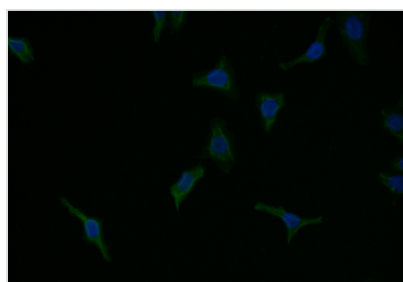




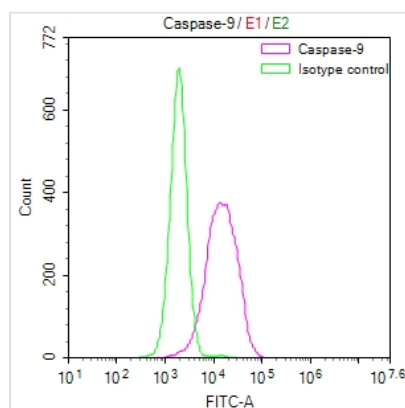
# CASP9 Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA940979A0HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P55211
<b>Immunogen</b>	A synthesized peptide derived from Human CASP9
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	ELISA, IF, FC; Recommended dilution: IF:1:50-1:200, 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>	Cancer;Cell biology;Metabolism
<b>Gene Names</b>	CASP9
<b>Clone No.</b>	8G5

## Image



Immunofluorescence staining of HepG2 with CSB-RA940979A0HU at 1:7, counter-stained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 525-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Overlay Peak curve showing HepG2 cells stained with CSB-RA940979A0HU (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 (1μg/1\*10<sup>6</sup>cells) for 45min at 4°C. The secondary antibody used was FITC-conjugated Goat Anti-rabbit IgG(H+L) at 1:200 dilution for 35min at 4°C. Control antibody (green line) was rabbit IgG (1μg/1\*10<sup>6</sup>cells) used under the same conditions. Acquisition of >10,000 events was performed.



## Description

The process of generating a recombinant monoclonal antibody against CASP9 began with the immunization of a rabbit using a synthesized peptide from human CASP9 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 was employed as a template to extend CASP9 antibody genes using degenerate primers. These extended CASP9 antibody genes were incorporated into a plasmid vector and transfected into host cells for expression. The CASP9 recombinant monoclonal antibody was then purified from the cell culture supernatant through affinity chromatography and subjected to ELISA, IF, and FC applications. It shows specific reactivity with human CASP9 protein.

CASP9 is a key regulator of apoptosis, serving as the initiator caspase in the intrinsic pathway. Its activation marks the commitment of a cell to undergo programmed cell death, a fundamental process in development, tissue homeostasis, and the elimination of damaged or potentially harmful cells.