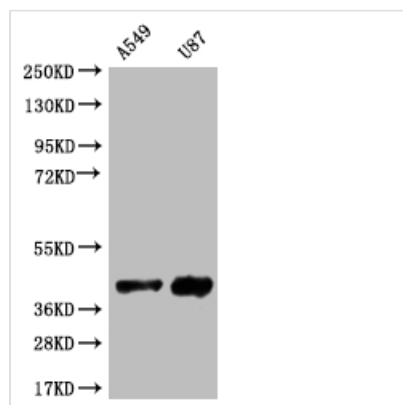




# SRR Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA945827A0HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	Q9GZT4
<b>Immunogen</b>	A synthesized peptide derived from human SRR
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	ELISA, WB, FC; Recommended dilution: WB:1:500-1:2000, FC:1:50-1:200
<b>Relevance</b>	Catalyzes the synthesis of D-serine from L-serine. D-serine is a key coagonist with glutamate at NMDA receptors. Has dehydratase activity towards both L-serine and D-serine. {ECO:0000269 PubMed:11054547, ECO:0000269 PubMed:20106978}.
<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>	Neuroscience
<b>Gene Names</b>	SRR
<b>Clone No.</b>	12B2

## Image



### Western Blot

Positive WB detected in: A549 whole cell lysate, U87 whole cell lysate

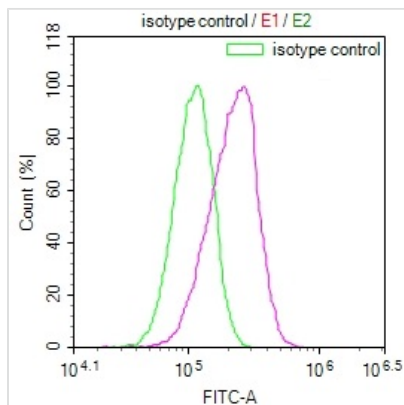
All lanes: SRR antibody at 1:1000

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 37 kDa

Observed band size: 37 kDa



Overlay Peak curve showing HepG2 cells stained with CSB-RA945827A0HU (red line) at 1:100. 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 (1ug/1\*10<sup>6</sup>cells) for 45min at 4?. The secondary antibody used was FITC-conjugated Goat Anti-rabbit IgG(H+L) at 1:200 dilution for 35min at 4?. Control antibody (green line) was rabbit IgG (1ug/1\*10<sup>6</sup>cells) used under the same conditions. Acquisition of >10,000 events was performed.

## Description

The SRR monoclonal antibody was generated using the synthesized peptide derived from human SRR protein as the immunogen. The gene sequence of the SRR monoclonal antibody was obtained by sequencing the cDNA and cloning it into a plasmid vector, which was then transfected into a suitable host cell. Following that, the resulting SRR recombinant monoclonal antibody was purified using affinity chromatography and its specificity was tested in ELISA, WB, and FC. It can specifically recognize human SRR protein.

Serine racemase (SRR) is an enzyme that catalyzes the conversion of L-serine to D-serine, an important co-agonist of the N-methyl-D-aspartate (NMDA) receptor, which plays a critical role in the regulation of synaptic plasticity, learning, and memory. D-serine acts by binding to the glycine site of the NMDA receptor and enhancing its activity. Serine racemase mainly regulates the levels of D-serine in the brain and modulate the activity of the NMDA receptor.