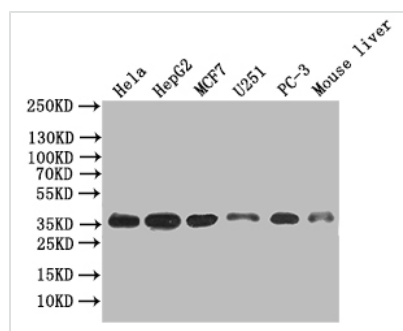




# ANXA4 Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA001845MA1HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P09525
<b>Immunogen</b>	Recombinant Human ANXA4 protein
<b>Species Reactivity</b>	Human, Mouse
<b>Tested Applications</b>	ELISA, WB, FC; Recommended dilution: WB:1:1000-1:5000, FC:1:20-1:200
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	Mouse IgG
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Recombinant Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Cardiovascular;Signal transduction
<b>Gene Names</b>	ANXA4
<b>Clone No.</b>	31D6

## Image



### Western Blot

Positive WB detected in: HeLa whole cell lysate, HepG2 whole cell lysate, MCF7 whole cell lysate, U251 whole cell lysate, PC-3 whole cell lysate, Mouse Liver tissue lysate

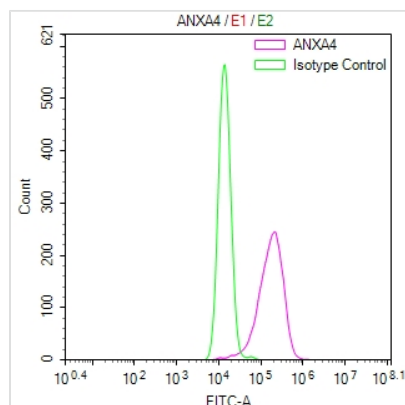
All lanes: ANXA4 antibody at 1:500

### Secondary

Goat polyclonal to mouse IgG at 1/50000 dilution

Predicted band size: 36 kDa

Observed band size: 36 kDa



Overlay Peak curve showing Hela cells stained with CSB-RA001845MA1HU (red line) at 1:800. The cells were fixed in 4% formaldehyde (15min) and permeated by 0.2% TritonX-100 for 10min. Then 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (1 $\mu$ g/1\*10<sup>6</sup> cells) for 45min at 4 $^{\circ}$ C. The secondary antibody used was FITC-conjugated Goat Anti-Mouse IgG(H+L) at 1/200 dilution for 35 min at 4 $^{\circ}$ C. Isotype control antibody (green line) was mouse IgG1 (1 $\mu$ g/1\*10<sup>6</sup> cells) used under the same conditions. Acquisition of >10,017 events was performed.