

Product Data Sheet

Anti-SerRS Antibody

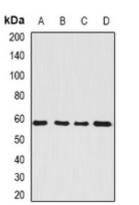
Catalog #	Source	Reactivity	Applications		
CQA1458	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Rab	bit polyclonal antibody	to SerRS		
Immunogen	Rec	combinant full length pro	otein of human SerRS		
Purification	The	antibody was purified	oy immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous lev	vels of SerRS protein.		
Clonality	Poly	yclonal			
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	1 0.01% sodium azide.			
Dilution	WB	(1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	SAF	۲S			
Alternative Na	ames SER	S; SerinetRNA ligase, d	cytoplasmic; Seryl-tRNA synthetase; SerRS;		
	Ser	yl-tRNA(Ser/Sec) synthe	tase		
Entrez Gene		6301 (Human); 20226 (Mouse); 266975 (Rat)			
SwissProt	P49	9591 (Human); P26638 (Mouse); Q6P799 (Rat)		
Storage/Stabi	l <mark>ity</mark> Ship	pped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	free	eze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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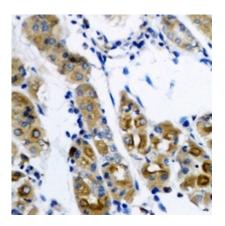




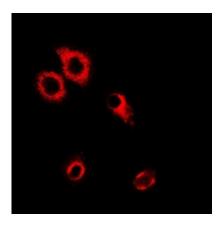
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Western blot analysis of SerRS expression in HepG2 (A), Hela (B), mouse kidney (C), rat brain (D) whole cell lysates. (Predicted band size: 58 kD; Observed band size: 59 kD)



Immunohistochemical analysis of SerRS staining in human gastric cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of SerRS staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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