

Product Data Sheet

Anti-IRS1 (pS307) Antibody

Catalog #	Source	Reactivity	Applications	
CPA4442	Rabbit	H, M, R, Mk	WB, IH, IF/IC	
Description	Ra	abbit polyclonal antibody to	o IRS1 (pS307)	
Immunogen	KI	LH-conjugated synthetic pe	otide encompassing a sequence within the center	
	re	gion of human IRS1 (pS307). The exact sequence is proprietary.	
Purification	Tł	ne antibody was purified by	immunogen affinity chromatography.	
Specificity	Re	ecognizes endogenous level	s of IRS1 (pS307) protein.	
Clonality	Po	olyclonal		
Form	Lie	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ar	nd 0.01% sodium azide.		
Dilution	Ŵ	/B (1/500 - 1/1000), IH (1/100) - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	IR	S1		
Alternative N	ames In	sulin receptor substrate 1;	IRS-1	
Entrez Gene	36	667 (Human); 16367 (Mous	e); 25467 (Rat)	
SwissProt	PS	35568 (Human); P35569 (M	ouse); P35570 (Rat)	
Storage/Stabi	i lity Sł	nipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
	fr	eeze/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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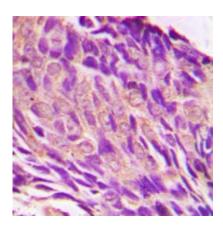
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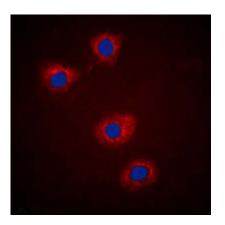
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Western blot analysis of IRS1 (pS307) expression in HEK293T (A), NS-1 (B) whole cell lysates.

Immunohistochemical analysis of IRS1 (pS307) staining in human prostate 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. w



Immunofluorescent analysis of IRS1 (pS307) staining in HEK293T 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. DAPI was used to stain the cell nuclei (blue).

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