

# **Product Data Sheet**

### Anti-IRS1 (Phospho-S312) Antibody

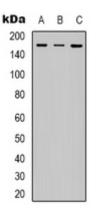
Catalog #	Source	Reactivity	Applications		
CPA5091	Rabbit	H, M, R	WB, IF/IC, IP		
Description	Ra	Rabbit polyclonal antibody to IRS1 (Phospho-S312)			
Immunogen	KL	H-conjugated synthetic ph	osphopeptide corresponding to residues surrounding		
	\$3	12 of human IRS1 protein	The exact sequence is proprietary.		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous leve	els of IRS1 protein only when phosphorylated at S312.		
Clonality	Ро	lyclonal			
Conjugation					
Form	Liq	ุ่นid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IF/IC (1/	100 - 1/500), IP (1/10 - 1/100)		
Gene Symbol	IRS	\$1			
Alternative N	ames Ins	sulin receptor substrate 1;	IRS-1		
Entrez Gene	36	67 (Human); 16367 (Mou	se); 25467 (Rat)		
SwissProt	Р3	5568 (Human); P35569 (N	1ouse); P35570 (Rat)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fre	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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Western blot analysis of IRS1 (Phospho-S312) expression in HepG2 (A), Hela (B), mouse spleen (C) whole cell lysates. (Predicted band size: 131 kD; Observed band size: 180 kD)



Immunofluorescent analysis of IRS1 (Phospho-S312) staining in HepG2 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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