

## **Product Data Sheet**

## Anti-Insulin Receptor (Phospho-Y1355) Antibody

Catalog #	Source	Reactivity	Applications		
CPA4440	Rabbit	H, M, R, Mk	WB, IH, IF/IC		
Description	o Insulin Receptor (Phospho-Y1355)				
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
	Y	1355 of human Insulin Reco	eptor protein. The exact sequence is proprietary.		
Purification The antibody was purified by immunogen affinity chromatography.					
Specificity Recognizes endogenous levels of Insulin Receptor protein only when phosp					
	at	t Y1355.			
Clonality	P	olyclonal			
Conjugation					
Form	Li	iquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	a	nd 0.01% sodium azide.			
Dilution	W	√B (1⁄500 - 1⁄1000), IH (1⁄50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	IN	NSR			
Alternative N	ames In	nsulin receptor; IR; CD220			
Entrez Gene	3	3643 (Human); 16337 (Mouse)			
SwissProt	P	06213 (Human); P15208 (N	1ouse); P15127 (Rat)		
Storage/Stabi	lity Sl	hipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fr	reeze/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

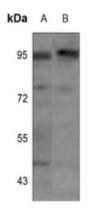
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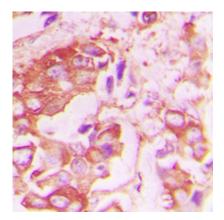
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For research purposes only, not for human use

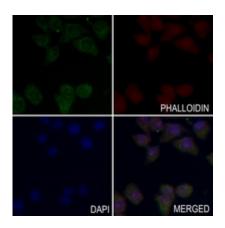
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Western blot analysis of Insulin Receptor (Phospho-Y1355) expression in mouse muscle (A), rat muscle (B) whole cell lysates. (Predicted band size: 156 kD; Observed band size: 95 kD)



Immunohistochemical analysis of Insulin Receptor (Phospho-Y1355) staining in human lung 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 Insulin Receptor (Phospho-Y1355) staining in LO2 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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