

Product Data Sheet

Anti-HGS Antibody

Catalog #	Source	Reactivity	Applications	
CPA3365	Rabbit	H, M, R, B, P	WB, IH, IF/IC	
Description	R	abbit polyclonal antibody to	HGS	
Immunogen	K	(LH-conjugated synthetic pep	tide encompassing a sequence within the center	
	re	egion of human HGS. The ex	act sequence is proprietary.	
Purification	Т	he antibody was purified by	immunogen affinity chromatography.	
Specificity	R	ecognizes endogenous level	s of HGS protein.	
Clonality	Р	Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	а	nd 0.01% sodium azide.		
Dilution	V	VB (1/500 - 1/1000), IH (1/100	- 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	Н	IGS		
Alternative Na	ames H	IRS; Hepatocyte growth facto	pr-regulated tyrosine kinase substrate; Hrs; Protein	
	р	pp110		
Entrez Gene	9	146 (Human); 15239 (Mouse	e); 56084 (Rat)	
SwissProt	C	014964 (Human); Q99LI8 (Mo	ouse); Q9JJ50 (Rat)	
Storage/Stabi	lity S	hipped at 4°C. Upon delivery	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

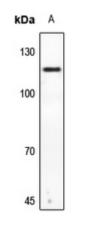
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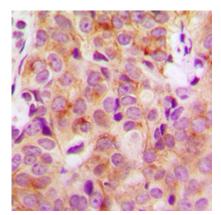
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For research purposes only, not for human use

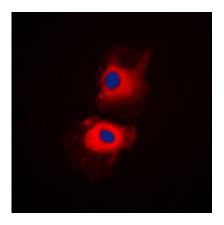
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Western blot analysis of HGS expression in Hela (A) whole cell lysates. (Predicted band size: 86 kD; Observed band size: 115 kD)



Immunohistochemical analysis of HGS staining in human breast 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 HGS staining in H1299 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 humidified 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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