

Product Data Sheet

Anti-GGH Antibody

Catalog #	Source	Reactivity	Applications		
CPA5212	Rabbit	H, M, R	WB, IH		
Description	Rab	bit polyclonal antibody	to GGH		
Immunogen	KLH	-conjugated synthetic _l	peptide encompassing a sequence within the C-term		
	regi	region of human GGH. The exact sequence is proprietary.			
Purification	The	antibody was purified	by immunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous le	vels of GGH protein.		
Clonality	Poly	rclonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000) <i>,</i> IH (1/5	0 - 1/100)		
Gene Symbol	GGH	ł			
Alternative N	ames Gan	ma-glutamyl hydrolas	e; Conjugase; GH; Gamma-Glu-X carboxypeptidase		
Entrez Gene	883	6 (Human); 14590 (Mo	use); 25455 (Rat)		
SwissProt	Q92	820 (Human); Q9Z0L8	(Mouse); Q62867 (Rat)		
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	free	ze/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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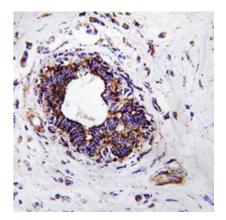
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Western blot analysis of GGH expression in mouse liver (A), mouse lung (B), rat liver (C), rat lung (D) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 36 kD)



Immunohistochemical analysis of GGH 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.

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