

Product Data Sheet

Anti-GALR1 Antibody

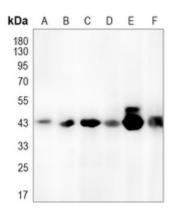
Catalog #	Source	Reactivity	Applications
CPA3612	Rabbit	H, M, R	WB, IF/IC
Description	Rab	bit polyclonal antibody t	o GALR1
Immunogen	KLH	I-conjugated synthetic pe	ptide encompassing a sequence within the center
	regi	ion of human GALR1. The	e exact sequence is proprietary.
Purification	The	e antibody was purified b	y immunogen affinity chromatography.
Specificity	Rec	ognizes endogenous leve	els of GALR1 protein.
Clonality	Poly	yclonal	
Conjugation			
Form	Liqu	uid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	l 0.01% sodium azide.	
Dilution	WB	(1/500 - 1/1000), IF/IC (1/50 - 1/200)
Gene Symbol	GAI	LR1	
Alternative Na	ames GAI	LNR; GALNR1; Galanin re	ceptor type 1; GAL1-R; GALR-1
Entrez Gene	258	37 (Human); 14427 (Mou	se); 50577 (Rat)
SwissProt	P47	7211 (Human); P56479 (N	1ouse); Q62805 (Rat)
Storage/Stabi	lity Ship	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	free	eze/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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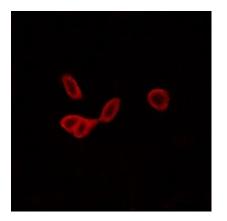




For research purposes only, not for human use

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Western blot analysis of GALR1 expression in LO2 (A), A2780 (B), THP1 (C), mouse brain (D), mouse liver (E), rat liver (F) whole cell lysates. (Predicted band size: 38 kD; Observed band size: 43 kD)



Immunofluorescent analysis of GALR1 staining in LOVO 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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