

Product Data Sheet

Anti-GPR87 Antibody

Catalog #	Source	Reactivity	Applications
CPA7349	Rabbit	H <i>,</i> M	WB, IF/IC
Description	Rat	obit polyclonal antibody	to GPR87
Immunogen	KLH	I-conjugated synthetic p	eptide encompassing a sequence within the center
	reg	ion of human GPR87. Th	e exact sequence is proprietary.
Purification	The	e antibody was purified l	by immunogen affinity chromatography.
Specificity	Rec	cognizes endogenous lev	els of GPR87 protein.
Clonality	Pol	yclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium ı	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WB	8 (1/500 - 1/1000), IF/IC (2	/100 - 1/500)
Gene Symbol	GP	R87	
Alternative Na	ames GPI	R95; G-protein coupled i	eceptor 87; G-protein coupled receptor 95
Entrez Gene	538	836 (Human); 84111 (Mo	buse)
SwissProt	Q9	BY21 (Human); Q99MT7	(Mouse)
Storage/Stabi	lity Shi	pped at 4°C. Upon delive	ery 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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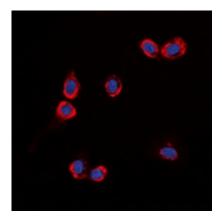
kDa
A
B
C
D
E

130
95
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For research purposes only, not for human use

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Western blot analysis of GPR87 expression in Myla2059 (A), PC3 (B), A549 (C), DLD (D), HepG2 (E) whole cell lysates. (Predicted band size: 41 kD; Observed band size: 41 kD)



Immunofluorescent analysis of GPR87 staining in HuvEc 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. DAPI was used to stain the cell nuclei (blue).

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