

## **Product Data Sheet**

## Anti-GPR172B Antibody

Catalog #	Source	Reactivity	Applications		
CPA4828	Rabbit	H, M, R	WB, IF/IC		
Description		Rabbit polyclonal antibody to GPR172B			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human GPR172B. T	he exact sequence is proprietary.		
Purification		The antibody was purified b	v immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of GPR172B protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid 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), IF/IC (1/	100 - 1/500)		
Gene Symbol		SLC52A1			
Alternative Na	ames	GPR172B; PAR2; RFT1; Solut	e carrier family 52, riboflavin transporter, member 1;		
		Porcine endogenous retrovi	us A receptor 2; PERV-A receptor 2; Protein GPR172B;		
		Riboflavin transporter 1; hRI	T1		
Entrez Gene		55065 (Human)			
SwissProt		Q9NWF4 (Human)			
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
		freeze/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

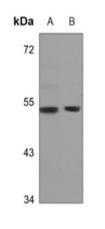
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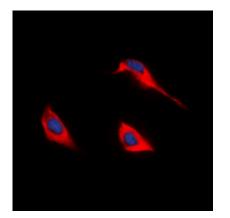
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Western blot analysis of GPR172B expression in mouse muscle (A), rat kidney (B) whole cell lysates. (Predicted band size: 46 kD; Observed band size: 46 kD)



Immunofluorescent analysis of GPR172B staining in HEK293T 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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