

Product Data Sheet

Anti-GPR119 Antibody

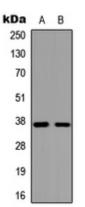
Catalog #	Source	Reactivity	Applications
CPA3970	Rabbit	H, Mk	WB, IH, IF/IC
Description	R	abbit polyclonal antibody	to GPR119
Immunogen	K	LH-conjugated synthetic p	eptide encompassing a sequence within the center
	re	egion of human GPR119. T	he exact sequence is proprietary.
Purification	TI	he antibody was purified	by immunogen affinity chromatography.
Specificity	R	ecognizes endogenous lev	els of GPR119 protein.
Clonality	Po	olyclonal	
Conjugation			
Form	Li	iquid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	aı	nd 0.01% sodium azide.	
Dilution	W	VB (1/500 - 1/1000), IH (1/1	00 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	G	PR119	
Alternative Na	a <mark>mes</mark> G	lucose-dependent insulin	otropic receptor; G-protein coupled receptor 119
Entrez Gene	13	39760 (Human); 236781 (Mouse); 302813 (Rat)
SwissProt	Q	8TDV5 (Human); Q7TQP3	(Mouse); Q7TQN8 (Rat)
Storage/Stabi	lity Sł	hipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid
	fr	eeze/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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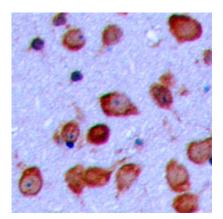
Cohesion



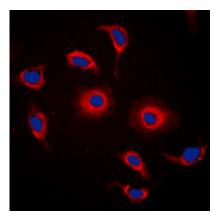
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Western blot analysis of GPR119 expression in HeLa (A), COLO205 (B) whole cell lysates. (Predicted band size: 36 kD; Observed band size: 37 kD)



Immunohistochemical analysis of GPR119 staining in human brain 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 GPR119 staining in HeLa 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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