

Product Data Sheet

Anti-GPR50 Antibody

| Catalog # | Source | Reactivity | Applications |
|---------------|----------|------------------------------|--|
| CPA2375 | Rabbit | H, M, R | WB, IF/IC |
| Description | Ral | bbit polyclonal antibody | to GPR50 |
| Immunogen | KLł | H-conjugated synthetic p | eptide encompassing a sequence within the center |
| | reg | gion of human GPR50. Th | e exact sequence is proprietary. |
| Purification | The | e antibody was purified b | oy immunogen affinity chromatography. |
| Specificity | Red | cognizes endogenous lev | els of GPR50 protein. |
| Clonality | Pol | lyclonal | |
| Conjugation | | | |
| Form | Liq | uid in 0.42% Potassium p | hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, |
| | and | d 0.01% sodium azide. | |
| Dilution | WE | 3 (1/500 - 1/1000), IF/IC (1 | /50 - 1/200) |
| Gene Symbol | GP | R50 | |
| Alternative N | ames Me | elatonin-related receptor | ; G protein-coupled receptor 50; H9 |
| Entrez Gene | 924 | 48 (Human); 14765 (Mou | ise) |
| SwissProt | Q1 | 3585 (Human); O88495 | Mouse); Q62953 (Rat) |
| Storage/Stabi | lity Shi | ipped at 4°C. Upon delive | ry aliquot and store at -20°C for one year. Avoid |
| | fre | 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

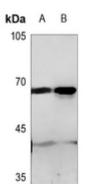
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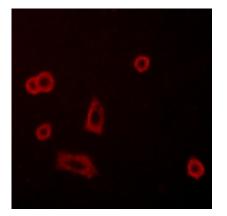


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Western blot analysis of GPR50 expression in mouse testis (A), rat testis (B) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 67 kD)



Immunofluorescent analysis of GPR50 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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