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GPR30 Rabbit pAb

Catalog Number: bs-1380R

Target Protein: GPR30
Concentration: 1mg/ml

Form: Liquid

Host: Rabbit

Clonality: Polyclonal

Isotype: IgG

Applications: WB (1:500-2000), IHC-P (1:100-500), IHC-F (1:100-500), IF (1:100-500), Flow-Cyt (1µg/Test),

ICC/IF (1:100)

Reactivity: Human, Mouse, Rat (predicted:Dog)

Predicted MW: 42 kDa Entrez Gene: 2852 Swiss Prot: Q99527

Source: KLH conjugated synthetic peptide derived from human GPR30: 251-375/375.

Purification: affinity purified by Protein A

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

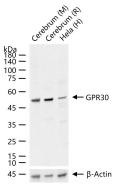
Background: This gene is a member of the G-protein coupled receptor 1 family and encodes a multi-pass

membrane protein that localizes to the endoplasmic reticulum. The protein binds estrogen, resulting in intracellular calcium mobilization and synthesis of phosphatidylinositol 3,4,5-trisphosphate in the nucleus. This protein therefore plays a role in the rapid nongenomic signaling events widely observed following stimulation of cells and tissues with estrogen.

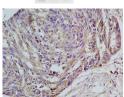
Alternate transcriptional splice variants which encode the same protein have been

characterized. [provided by RefSeq, Jul 2008]

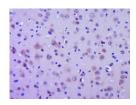
VALIDATION IMAGES



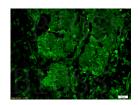
25 ug total protein per lane of various lysates (see on figure) probed with GPR30 polyclonal antibody, unconjugated (bs-1380R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



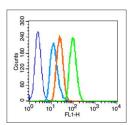
Tissue/cell: human rectal carcinoma; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-GPR30 Polyclonal Antibody, Unconjugated (bs-1380R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



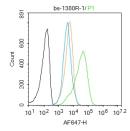
Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (GPR30) Polyclonal Antibody, Unconjugated (bs-1380R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Tissue/cell: human colon carcinoma;4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-GPR30 Polyclonal Antibody, Unconjugated(bs-R) 1:200, overnight at 4°C; The secondary antibody was Goat Anti-Rabbit IgG, Cy3 conjugated(bs-0295G-Cy3)used at 1:200 dilution for 40 minutes at 37°C.



Blank control (blue line): A431 cells (blue). Primary Antibody (green line): Rabbit Anti-GPR30 antibody (bs-1380R) Dilution: $1\mu g/10^6$ cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-FITC Dilution: $1\mu g$ /test. Protocol The cells were fixed with 70% methanol (Overnight at 4°C) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C. Cells stained with Primary Antibody for 30 min at room temperature. The cells were then incubated in 1 X PBS/2%BSA/10% goat serum to block non-specific protein-protein interactions followed by the antibody for 15 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.



Blank control: HepG2. Primary Antibody (green line): Rabbit Anti-GPR30 antibody (bs-1380R) Dilution: $1\mu g / 10^6$ cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody: Goat anti-rabbit IgG-AF647 Dilution: $1\mu g / \text{test}$. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 0.1% PBST for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature . Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

PRODUCT SPECIFIC PUBLICATIONS

[IF=7.129] Qianfeng Liu. et al. Perfluoroalkyl substances promote breast cancer progression via ER α and GPER mediated PI3K/Akt and MAPK/Erk signaling pathways. ECOTOX ENVIRON SAFE. 2023 Jun;258:114980 WB; Human . 37148752

[IF=5.6] Zihao Fang. et al. 17β-Estradiol mediates TGFBR3/Smad2/3 signaling to attenuate the fibrosis of TGF-β1-induced bovine endometrial epithelial cells via GPER. J CELL PHYSIOL. 2023 Nov;: WB,IHC; Bovine . 37991438

[IF=4.8] Lixia Chen. et al. Activation of GPER1 by G1 prevents PTSD-like behaviors in mice: Illustrating the mechanisms from BDNF/TrkB to mitochondria and synaptic connection. CNS NEUROSCI THER. 2024 Jul;30(7):e14855 WB; MOUSE. 38992889

[IF=3.75] Andressa S. Sousa. et al. Evaluation of coronary function in female rats with severe type 1 diabetes: Effects of combined treatment with insulin and pyridoxamine. MICROVASC RES. 2022 Dec;:104474 WB; Rat . 36592817

[IF=3.889] Ting Chen. et al. Tongmai Yangxin pill alleviates myocardial no-reflow by activating GPER to regulate HIF-1 α signaling and downstream potassium channels. PHARM BIOL. 2023;61(1):499-513 WB; Rat . 36896463