

mPR γ (C-20): sc-28021

BACKGROUND

The steroid progesterone induces the resumption of maturation in oocytes via a nongenomic pathway through binding to a novel, membrane progesterin receptor (mPR). This pathway inhibits adenylyl cyclase and reduces intracellular cAMP, and also activates mitogen-activated protein kinase to effect signal transduction pathways. Five distinct groups, designated α , β , γ , δ and ϵ , comprise this gene family, and while all contain 7 transmembrane domains, they show distinct distributions in reproductive, neural, kidney and intestinal tissues, respectively. These characteristics separate them from nuclear progesterin receptors, and instead imply similarity to G-protein coupled receptors.

REFERENCES

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- Zhu, Y., et al. 2003. Cloning, expression, and characterization of a membrane progesterin receptor and evidence it is an intermediary in meiotic maturation of fish oocytes. *Proc. Natl. Acad. Sci. USA* 100: 2231-2236.
- Kudwa, A.E., et al. 2003. Double oestrogen receptor α and β knockout mice reveal differences in neural oestrogen-mediated progesterin receptor induction and female sexual behaviour. *J. Neuroendocrinol.* 15: 978-983.
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- Kudwa, A.E., et al. 2004. Estrogen receptor β modulates estradiol induction of progesterin receptor immunoreactivity in male, but not in female, mouse medial preoptic area. *Endocrinology* 145: 4500-4506.
- Lonstein, J.S., et al. 2004. Immunocytochemical investigation of nuclear progesterin receptor expression within dopaminergic neurones of the female rat brain. *J. Neuroendocrinol.* 16: 534-543.

CHROMOSOMAL LOCATION

Genetic locus: PAQR5 (human) mapping to 15q23.

SOURCE

mPR γ (C-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of membrane progesterin receptor gamma of human origin.

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-28021 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

RESEARCH USE

For research use only, not for use in diagnostic procedures.

APPLICATIONS

mPR γ (C-20) is recommended for detection of mPR γ of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

mPR γ (C-20) is also recommended for detection of mPR γ in additional species, including equine, canine, bovine and porcine.

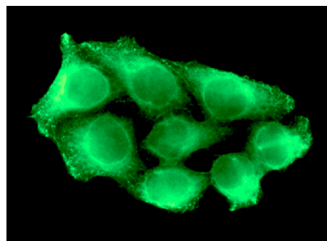
Suitable for use as control antibody for mPR γ siRNA (h): sc-106235, mPR γ shRNA Plasmid (h): sc-106235-SH and mPR γ shRNA (h) Lentiviral Particles: sc-106235-V.

Molecular Weight of mPR γ : 38 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker[™] compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker[™] Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz[™] Mounting Medium: sc-24941.

DATA



mPR γ (C-20): sc-28021. Immunofluorescence staining of methanol-fixed HeLa cells showing membrane localization.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

MONOS
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Try mPR δ/γ (B-8): **sc-514273**, our highly recommended monoclonal alternative to mPR γ (C-20).