

mPR δ / γ (E-17): sc-28020

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 adenyl 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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- Curran-Rauhut, M.A., et al. 2002. The distribution of progesterin receptor mRNA in rat brainstem. *Brain Res. Gene Expr. Patterns* 1: 151-157.
- 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.
- Zhu, Y., et al. 2003. Identification, classification, and partial characterization of genes in humans and other vertebrates homologous to a fish membrane progesterin receptor. *Proc. Natl. Acad. Sci. USA* 100: 2237-2242.
- 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.
- 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.

CHROMOSOMAL LOCATION

Genetic locus: PAQR5 (human) mapping to 15q23, PAQR6 (human) mapping to 1q22.

SOURCE

mPR δ / γ (E-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of mPR γ 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-28020 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

STORAGE

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

APPLICATIONS

mPR δ / γ (E-17) is recommended for detection of mPR γ and mPR δ of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], 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); may cross-react with mPR ϵ and PAQR4.

mPR δ / γ (E-17) is also recommended for detection of mPR γ and mPR δ in additional species, including equine, canine, bovine and porcine.

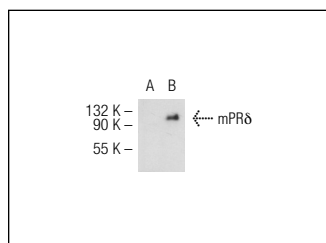
Molecular Weight of mPR δ / γ : 38 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200 or mPR δ (h): 293T Lysate: sc-115979.

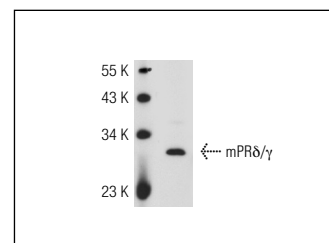
RECOMMENDED SECONDARY REAGENTS

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) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) 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 δ / γ (E-17): sc-28020. Western blot analysis of mPR δ expression in non-transfected: sc-117752 (A) and human mPR δ transfected: sc-115979 (B) 293T whole cell lysates.



mPR δ / γ (E-17): sc-28020. Western blot analysis of mPR δ / γ expression in HeLa whole cell lysate.

RESEARCH USE

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

PROTOCOLS

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