

mPR γ (H-70): sc-292399

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

- Sheng, Y., et al. 2001. Regulation of *Xenopus* oocyte meiosis arrest by G protein $\beta\gamma$ subunits. *Curr. Biol.* 11: 405-416.
- Curran-Rauhut, M.A., et al. 2002. The distribution of progesterin receptor mRNA in rat brainstem. *Brain Res. Gene Expr. Patterns* 1: 151-157.

CHROMOSOMAL LOCATION

Genetic locus: PAQR5 (human) mapping to 15q23; Paqr5 (mouse) mapping to 9 B.

SOURCE

mPR γ (H-70) is a rabbit polyclonal antibody raised against amino acids 261-330 mapping at 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. Also available as TransCruz reagent for Gel Supershift and ChIP applications, sc-292399 X, 200 μ g/0.1 ml.

APPLICATIONS

mPR γ (H-70) is recommended for detection of mPR γ of mouse, rat and 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).

mPR γ (H-70) is also recommended for detection of mPR γ in additional species, including equine, bovine, porcine and canine.

Suitable for use as control antibody for mPR γ siRNA (h): sc-106235, mPR γ siRNA (m): sc-155918, mPR γ shRNA Plasmid (h): sc-106235-SH, mPR γ shRNA Plasmid (m): sc-155918-SH, mPR γ shRNA (h) Lentiviral Particles: sc-106235-V and mPR γ shRNA (m) Lentiviral Particles: sc-155918-V.

mPR γ (H-70) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

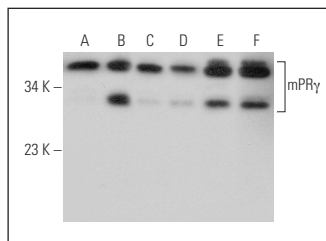
Molecular Weight of mPR γ : 38 kDa.

Positive Controls: HEK293 whole cell lysate: sc-45136, HUV-EC-C whole cell lysate: sc-364180 or Jurkat whole cell lysate: sc-2204.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (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 goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



mPR γ (H-70): sc-292399. Western blot analysis of mPR γ expression in Jurkat (A), HeLa (B), SW480 (C), PC-12 (D), HUV-EC-C (E) and HEK293 (F) whole cell lysates.

STORAGE

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

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.

MONOS
Satisfaction
Guaranteed

Try mPR δ/γ (B-8): sc-514273, our highly recommended monoclonal alternative to mPR γ (H-70).