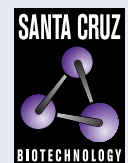


mPR $\delta/\gamma$  (B-8): sc-514273

The Power to Question

## 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  $\alpha$ ,  $\beta$ ,  $\gamma$ ,  $\delta$  and  $\epsilon$ , 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.
- 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  $\alpha$  and  $\beta$  knockout mice reveal differences in neural oestrogen-mediated progesterin receptor induction and female sexual behaviour. *J. Neuroendocrinol.* 15: 978-983.

## CHROMOSOMAL LOCATION

Genetic locus: PAQR6 (human) mapping to 1q22, PAQR5 (human) mapping to 15q23; Paqr6 (mouse) mapping to 3 F1, Paqr5 (mouse) mapping to 9 B.

## SOURCE

mPR $\delta/\gamma$  (B-8) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 234-257 near the C-terminus of mPR $\gamma$  of human origin.

## PRODUCT

Each vial contains 200  $\mu$ g IgG $_1$  kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

mPR $\delta/\gamma$  (B-8) is available conjugated to agarose (sc-514273 AC), 500  $\mu$ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-514273 HRP), 200  $\mu$ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-514273 PE), fluorescein (sc-514273 FITC), Alexa Fluor<sup>®</sup> 488 (sc-514273 AF488), Alexa Fluor<sup>®</sup> 546 (sc-514273 AF546), Alexa Fluor<sup>®</sup> 594 (sc-514273 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-514273 AF647), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-514273 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-514273 AF790), 200  $\mu$ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-514273 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

## APPLICATIONS

mPR $\delta/\gamma$  (B-8) is recommended for detection of mPR $\delta$  and mPR $\gamma$  of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ 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).

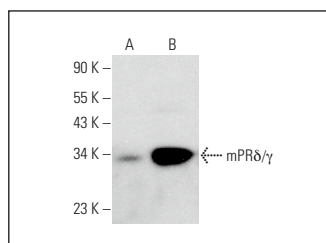
Molecular Weight of mPR $\delta/\gamma$ : 38 kDa.

Positive Controls: HUV-EC-C whole cell lysate: sc-364180, EOC 20 whole cell lysate: sc-364187 or WI-38 whole cell lysate: sc-364260.

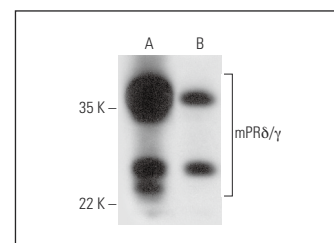
## RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>™</sup> Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 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 m-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz<sup>®</sup> Mounting Medium: sc-24941 or UltraCruz<sup>®</sup> Hard-set Mounting Medium: sc-359850.

## DATA



mPR $\delta/\gamma$  (B-8): sc-514273. Western blot analysis of mPR $\delta/\gamma$  expression in C6 (A) and EOC 20 (B) whole cell lysates.



mPR $\delta/\gamma$  (B-8): sc-514273. Western blot analysis of mPR $\delta/\gamma$  expression in HUV-EC-C (A) and WI-38 (B) 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](http://www.scbt.com) for detailed protocols and support products.

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