

Adenosine A2B-R (N-19): sc-7506

BACKGROUND

Adenosine is involved in a variety of processes, including the synthesis of urea, the anti-inflammatory response, and the inhibition of protein synthesis. The Adenosine receptors, including Adenosine A1-R, Adenosine A2A-R, Adenosine A2B-R and Adenosine A3-R, are integral membrane proteins that are members of the G protein-coupled receptor family. Adenosine A1-R mediates ureagenesis in a partially calcium-dependent manner. Adenosine is known to mediate coronary vasodilation via Adenosine A2A-R. Collagen synthesis and total protein synthesis are inhibited in certain cells by Adenosine, acting via the A2B receptors. Activation of Adenosine A3-R inhibits the induction of TNF α and blocks the endotoxin CD14 receptor signal transduction pathway.

REFERENCES

1. Mahan, L.C., et al. 1991. Cloning and expression of an A1 adenosine receptor from rat brain. *Mol. Pharmacol.* 40: 1-7.
2. Furlong, T.J., et al. 1992. Molecular characterization of a human brain adenosine A2 receptor. *Brain Res. Mol. Brain Res.* 15: 62-66.
3. Pierce, K.D., et al. 1992. Molecular cloning and expression of an adenosine A2B receptor from human brain. *Biochem. Biophys. Res. Commun.* 187: 86-93.
4. Salvatore, C.A., et al. 1993. Molecular cloning and characterization of the human A3 adenosine receptor. *Proc. Natl. Acad. Sci. USA* 90: 10365-10369.
5. McWhinney, C.D., et al. 1996. Activation of adenosine A3 receptors on macrophages inhibits tumor necrosis factor- α . *Eur. J. Pharmacol.* 310: 209-216.

CHROMOSOMAL LOCATION

Genetic locus: ADORA2B (human) mapping to 17p12; Adora2b (mouse) mapping to 11 B2.

SOURCE

Adenosine A2B-R (N-19) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the N-terminus of Adenosine A2B-R 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-7506 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.

RESEARCH USE

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

APPLICATIONS

Adenosine A2B-R (N-19) is recommended for detection of Adenosine A2B-R 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).

Adenosine A2B-R (N-19) is also recommended for detection of Adenosine A2B-R in additional species, including equine and canine.

Suitable for use as control antibody for Adenosine A2B-R siRNA (h): sc-29642, Adenosine A2B-R siRNA (m): sc-29643, Adenosine A2B-R shRNA Plasmid (h): sc-29642-SH, Adenosine A2B-R shRNA Plasmid (m): sc-29643-SH, Adenosine A2B-R shRNA (h) Lentiviral Particles: sc-29642-V and Adenosine A2B-R shRNA (m) Lentiviral Particles: sc-29643-V.

Molecular Weight (predicted) of Adenosine A2B-R: 36 kDa.

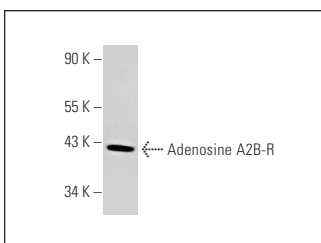
Molecular Weight (observed) of Adenosine A2B-R: 45 kDa.

Positive Controls: HT-29 whole cell lysate: sc-364232.

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



Adenosine A2B-R (N-19): sc-7506. Western blot analysis of Adenosine A2B-R expression in HT-29 whole cell lysate.

SELECT PRODUCT CITATIONS

1. Nguyen, D.K., et al. 2003. Th1 cytokines regulate adenosine receptors and their downstream signaling elements in human microvascular endothelial cells. *J. Immunol.* 171: 3991-3998.
2. Desrosiers, M.D., et al. 2007. Adenosine deamination sustains dendritic cell activation in inflammation. *J. Immunol.* 179: 1884-1892.