# P2Y5 (H-60): sc-20126



The Power to Question

## **BACKGROUND**

Nucleotides are emerging as important extracellular signaling molecules that mediate several effects, such as proliferation, differentiation, chemotaxis and cytokine release. The P2 receptor family is activated by the binding of nucleotides and is divided into two subfamilies, P2X and P2Y. The P2X receptor family is comprised of ligand-gated ion channels that allow for the increased permeability of calcium into the cell in response to extracellular ATP. The P2Y receptor family consists of eleven members of G protein-coupled receptors, P2Y1 through P2Y11, which mediate the effects of extracellular nucleotides, primarily through the activation of phospholipase C. To some extent, the P2Y receptors can also activate potassium channels or, alternatively, inhibit adenylate cyclase and N-type calcium channels in response to extracellular nucleotides. The P2Y receptors are differentially expressed in several tissue in leukocytes, which suggests a role for the P2Y receptor family in the activation of leukocytes and platelets in response to inflammation or vascular damage.

## **REFERENCES**

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- 2. North, R.A., et al. 1997. Nucleotide receptors. Curr. Opin. Neurobiol. 7: 346-357.
- Burnstock, G., et al. 2000. P2X receptors in sensory neurones. Br. J. Anaesth. 84: 476-488.
- Oury, C., et al. 2000. A natural dominant negative P2X1 receptor due to deletion of a single amino acid residue. J. Biol. Chem. 275: 22611-22614.
- Ding, S., et al. 2000. Inactivation of P2X2 purinoceptors by divalent cations. J. Physiol. 2: 199-214.
- Adrian, K., et al. 2000. Expression of purinergic receptors (ionotropic P2X1-7 and metabotropic P2Y1-11) during myeloid differentiation of HL60 cells. Biochim. Biophys. Acta 1492: 127-138.
- 7. Di Virgilio, F., et al. 2001. Nucleotide receptors: an emerging family of regulatory molecules in blood cells. Blood 97: 587-600.

# CHROMOSOMAL LOCATION

Genetic locus: LPAR6 (human) mapping to 13q14.2.

## SOURCE

P2Y5 (H-60) is a rabbit polyclonal antibody raised against amino acids 121-180 mapping within an internal region of P2Y5 of human origin.

## **PRODUCT**

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

# **STORAGE**

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

#### **APPLICATIONS**

P2Y5 (H-60) is recommended for detection of P2Y5 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg 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).

P2Y5 (H-60) is also recommended for detection of P2Y5 in additional species, including equine and porcine.

Suitable for use as control antibody for P2Y5 siRNA (h): sc-42583, P2Y5 shRNA Plasmid (h): sc-42583-SH and P2Y5 shRNA (h) Lentiviral Particles: sc-42583-V.

Molecular Weight (predicted) of P2Y5: 39 kDa.

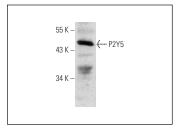
Molecular Weight (observed) of P2Y5: 46 kDa.

Positive Controls: Hs 732.Sk/Mu whole cell lysate: sc-364362.

## **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**



P2Y5 (H-60): sc-20126. Western blot analysis of P2Y5 expression in Hs 732.Sk/Mu whole cell lysate.

# **SELECT PRODUCT CITATIONS**

 Vila, A.M., et al. 2010. Development of a new magnetic beads-based immunoprecipitation strategy for proteomics analysis. J. Proteomics 73: 1491-1501.

## **RESEARCH USE**

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

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