# SANTA CRUZ BIOTECHNOLOGY, INC.

# PAR-4 (N-20): sc-21048



#### BACKGROUND

Thrombin receptor (also designated protease-activated receptor-1 or PAR-1), PAR-2, PAR-3 and PAR-4 compose a distinct class of G protein-coupled receptors activated by proteolysis. Cleavage of these receptors by proteases occurs within the amino-terminal extracellular domain. Thrombin, a serine protease involved in platelet aggregation and blood coagulation, activates the thrombin receptor, resulting in elevated intracellular calcium levels in platelets. Thrombin also cleaves PAR-3 *in vitro*, suggesting that PAR-3 may be involved in thrombosis or mitogenesis. Thrombin receptor and PAR-4 appear to account for most thrombin signaling in platelets. Activation of PAR-2 *in vitro* is induced by trypsin, suggesting that PAR-2 is not an alternative thrombin receptor. Cytokines including TNF- $\alpha$  and IL-1 $\beta$  increase PAR-2 expression, indicating PAR-2 involvement in the acute inflammatory response.

# REFERENCES

- Santulli, R.J., et al. 1995. Evidence for the presence of a protease-activated receptor distinct from the thrombin receptor in human keratinocytes. Proc. Natl. Acad. Sci. USA 92: 9151-9155.
- Lerner, D.J., et al. 1996. Agonist recognition by proteinase-activated receptor 2 and thrombin receptor. Importance of extracellular loop interactions for receptor function. J. Biol. Chem. 271: 13943-13947.
- 3. Nystedt, S., et al. 1996. The proteinase-activated receptor 2 is induced by inflammatory mediators in human endothelial cells. Comparison with the thrombin receptor. J. Biol. Chem. 271: 14910-14915.
- 4. Goldsack, N.R., et al. 1998. Thrombin. Int. J. Biochem. Cell Biol. 30: 641-646.
- 5. Xu, W.F., et al. 1998. Cloning and characterization of human proteaseactivated receptor 4. Proc. Natl. Acad. Sci. USA 95: 6642-6646.
- Sullivan, R., et al. 1998. Analysis of a Ca<sup>2+</sup>-activated K<sup>+</sup> channel that mediates hyperpolarization via the thrombin receptor pathway. Am. J. Physiol. 275: C1342-C1348.
- Schmidt, V.A., et al. 1998. The human proteinase-activated receptor-3 (PAR-3) gene. Identification within a Par gene cluster and characterization in vascular endothelial cells and platelets. J. Biol. Chem. 273: 15061-15068.
- Kahn, M.L., et al. 1999. Protease-activated receptors 1 and 4 mediate activation of human platelets by thrombin. J. Clin. Invest. 103: 879-887.

## CHROMOSOMAL LOCATION

Genetic locus: F2RL3 (human) mapping to 19p13.11; F2rl3 (mouse) mapping to 8 B3.3.

#### SOURCE

PAR-4 (N-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of PAR-4 of human origin.

# **STORAGE**

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

## PRODUCT

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

Blocking peptide available for competition studies, sc-21048 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

# **APPLICATIONS**

PAR-4 (N-20) is recommended for detection of PAR-4 of human and, to a lesser extent, mouse and rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), 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).

PAR-4 (N-20) is also recommended for detection of PAR-4 in additional species, including equine and porcine.

Suitable for use as control antibody for PAR-4 siRNA (h): sc-72068, PAR-4 siRNA (m): sc-72069, PAR-4 shRNA Plasmid (h): sc-72068-SH, PAR-4 shRNA Plasmid (m): sc-72069-SH, PAR-4 shRNA (h) Lentiviral Particles: sc-72068-V and PAR-4 shRNA (m) Lentiviral Particles: sc-72069-V.

Molecular Weight of PAR-4: 38 kDa.

Positive Controls: HL-60 whole cell lysate: sc-2209, Daudi cell lysate: sc-2415 or AML-193 whole cell lysate: sc-364182.

#### **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) Immunofluo-rescence: use donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

#### **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 Satisfation Guaranteed

Try **PAR-4 (5F4): sc-293206**, our highly recommended monoclonal alternative to PAR-4 (N-20).