# CysLT<sub>2</sub> Receptor (V-15): sc-27915



The Power to Question

#### **BACKGROUND**

Cysteinyl leukotriene (CysLTs) induce intracellular calcium mobilization through the binding of two distinct seven-transmembrane, G protein-coupled receptors, designated  $\mbox{CysLT}_1$  and  $\mbox{CysLT}_2$  Receptors, to induce potent bronchoconstriction. Airway smooth muscle and macrophages express both receptor types, and additionally monocytes and eosinophils express  $\mbox{CysLT}_1$  Receptor, while cardiac Purkinje cells, adrenal medulla, peripheral blood leukocytes and brain also utilize  $\mbox{CysLT}_2$  Receptor. The effects of the CysLT receptors can be blocked by antagonists, indicating a therapeutic mechanism for the treatment of asthma and allergies.

## **REFERENCES**

- Sarau, H.M., et al. 1999. Identification, molecular cloning, expression, and characterization of a cysteinyl leukotriene receptor. Mol. Pharmacol. 56: 657-663.
- Lynch, K.R., et al. 1999. Characterization of the human cysteinyl leukotriene CysLT<sub>1</sub> receptor. Nature 399: 789-793.
- 3. Heise, C.E., et al. 2000. Characterization of the human cysteinyl leukotriene 2 receptor. J. Biol. Chem. 275: 30531-30536.
- 4. Sjostrom, M., et al. 2001. Human umbilical vein endothelial cells generate leukotriene C4 via microsomal glutathione S-transferase type 2 and express the CysLT<sub>1</sub> receptor. Eur. J. Biochem. 268: 2578-2586.
- Maekawa, A., et al. 2001. Identification in mice of two isoforms of the cysteinyl leukotriene 1 receptor that result from alternative splicing. Proc. Natl. Acad. Sci. USA 98: 2256-2261.
- Leff, A.R. 2001. Regulation of leukotrienes in the management of asthma: biology and clinical therapy. Annu. Rev. Med. 52: 1-14.

# CHROMOSOMAL LOCATION

Genetic locus: CYSLTR2 (human) mapping to 13q14.2.

# **SOURCE**

CysLT<sub>2</sub> Receptor (V-15) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an N-terminus cytoplasmic domain of Cysteinyl leukotriene receptor 2 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.

Blocking peptide available for competition studies, sc-27915 P, (100  $\mu$ 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.

# **PROTOCOLS**

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

#### **APPLICATIONS**

CysLT<sub>2</sub> Receptor (V-15) is recommended for detection of CysLT<sub>2</sub> Receptor of human 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).

Suitable for use as control antibody for  $CysLT_2$  Receptor siRNA (h): sc-43713,  $CysLT_2$  Receptor shRNA Plasmid (h): sc-43713-SH and  $CysLT_2$  Receptor shRNA (h) Lentiviral Particles: sc-43713-V.

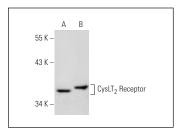
Molecular Weight of CysLT<sub>2</sub>: 43 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200, Jurkat whole cell lysate: sc-2204 or JAR cell lysate: sc-2276.

#### **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) 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**



<code>CysLT2</code> Receptor (V-15): sc-27915. Western blot analysis of <code>CysLT2</code> Receptor expression in HeLa ( $\bf A$  and JAR ( $\bf B$ ) whole cell lysates.

## **RESEARCH USE**

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

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