# OSMR β (AN-U2): sc-21797

### **BACKGROUND**

Oncostatin M (OSM) is a glycoprotein that inhibits the growth of a broad range of human tumor cell lines, but does not influence the growth of normal human fibroblasts. Expression of OSM is greatest in activated monocytic and lymphocytic cell lines and in normal adherent macrophages. Amino acid sequence analysis of OSM has revealed homology with leukemia inhibitory factor (LIF), granulocyte colony stimulating factor (G-CSF) and interleukin 6 (IL-6), all of which affect the growth and differentiation of a broad range of cell types, including those of hematopoietic origin. OSMR  $\beta$  (oncostatin M receptor  $\beta$ ), also known as OSMR, is a 979 amino acid single-pass type I membrane protein that functions as a receptor for OSM. Expressed at high levels in neural cells, as well as fibroblast and epithelial tumor lines, OSMR  $\beta$  exists as a hetero-dimer that interacts with interleukins and is able to transduce OSM-induced signaling events. Defects in the gene encoding OSMR  $\beta$  are the cause of primary cutaneous amyloidosis (PCA), an autosomal dominant disorder characterized by chronic itching of the skin.

### **REFERENCES**

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- 2. Blanchard, F., et al. 2001. Oncostatin M regulates the synthesis and turnover of gp130, leukemia inhibitory factor receptor  $\alpha$ , and oncostatin M receptor  $\beta$  by distinct mechanisms. J. Biol. Chem. 276: 47038-47045.
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- Radtke, S., et al. 2002. Novel role of Janus kinase 1 in the regulation of oncostatin M receptor surface expression. J. Biol. Chem. 277: 11297-11305.
- Dillon, S.R., et al. 2004. Interleukin 31, a cytokine produced by activated T cells, induces dermatitis in mice. Nat. Immunol. 5: 752-760.
- Abir, R., et al. 2005. Immunocytochemical detection and reverse transcription polymerase chain reaction expression of oncostatin M (OSM) and its receptor (OSMRβ) in human fetal and adult ovaries. Fertil. Steril. 83: 1188-1196.

### **CHROMOSOMAL LOCATION**

Genetic locus: OSMR (human) mapping to 5p13.1; Osmr (mouse) mapping to 15 A1.

## SOURCE

OSMR  $\beta$  (AN-U2) is a mouse monoclonal antibody raised against OSMR $\beta$ -transfected COS cells of human origin.

#### **RESEARCH USE**

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

### **PRODUCT**

Each vial contains 200  $\mu g$   $lgG_1$  kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

OSMR  $\beta$  (AN-U2) is available conjugated to either phycoerythrin (sc-21797 PE) or fluorescein (sc-21797 FITC), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM.

### **APPLICATIONS**

OSMR  $\beta$  (AN-U2) is recommended for detection of OSMR  $\beta$  of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)] and flow cytometry (1  $\mu$ g per 1 x 10<sup>6</sup> cells).

Suitable for use as control antibody for OSMR  $\beta$  siRNA (h): sc-40068, OSMR  $\beta$  siRNA (m): sc-40069, OSMR  $\beta$  shRNA Plasmid (h): sc-40068-SH, OSMR  $\beta$  shRNA Plasmid (m): sc-40069-SH, OSMR  $\beta$  shRNA (h) Lentiviral Particles: sc-40068-V and OSMR  $\beta$  shRNA (m) Lentiviral Particles: sc-40069-V.

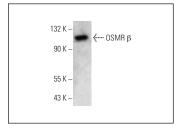
Molecular Weight of OSMR β: 180 kDa.

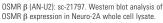
Positive Controls: Neuro-2A whole cell lysate: sc-364185 or HeLa whole cell lysate: sc-2200.

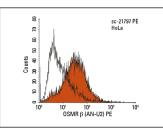
### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG $\kappa$  BP-HRP: sc-516102 or m-lgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</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).

### **DATA**







OSMR $\beta$  (AN-U2) PE: sc-21797 PE. FCM analysis of HeLa cells. Black line histogram represents the isotype control, normal mouse  $\lg G_1$ -PE: sc-2866.

## **SELECT PRODUCT CITATIONS**

 Hoermann, G., et al. 2012. Identification of oncostatin M as a JAK2 V617F-dependent amplifier of cytokine production and bone marrow remodeling in myeloproliferative neoplasms. FASEB J. 26: 894-906.

#### **STORAGE**

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