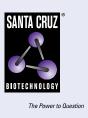
# SANTA CRUZ BIOTECHNOLOGY, INC.

# VPAC1 (B-4): sc-377152



# BACKGROUND

The vasoactive intestinal peptide (VIP) and pituitary adenylate cylaseactivating polypeptide (PACAP) belong to a superfamily of peptide hormones that include glucagon, secretin and growth hormone releasing hormone. The effects of VIP and PACAP are mediated by three G protein-coupled receptors, VPAC1, VPAC2 and the PACAP receptor (also designated PAC1-R). The VPAC receptors have equal affinities for VIP and PACAP, which stimulate the activation of adenylyl cyclase. Both VPAC1 and VPAC2 are abundantly expressed in brain and T cells, where they modulate neuronal differentiation and T cell activation, respectively The PACAP receptor is a seven transmembrane protein that produces at least eight isoforms by alternative splicing. Each isoform is associated with a specific signaling pathway and a specific expression pattern. The PACAP receptor, which is thought to play an integral role in brain development, preferentially binds PACAP in order to stimulate a cAMP-protein kinase A signaling pathway.

# REFERENCES

- Shen, S., et al. 2000. Overexpression of the human VPAC2 receptor in the suprachiasmatic nucleus alters the circadian phenotype of mice. Proc. Natl. Acad. Sci. USA 97: 11575-11580.
- Shioda, S. 2000. Pituitary adenylate cyclase-activating polypeptide (PACAP) and its receptors in the brain. Kaibogaku Zasshi 75: 487-507.
- Bajo, A.M., et al. 2000. Expression of vasoactive intestinal peptide (VIP) receptors in human uterus. Peptides 21: 1383-1388.

#### **CHROMOSOMAL LOCATION**

Genetic locus: VIPR1 (human) mapping to 3p22.1.

# SOURCE

VPAC1 (B-4) is a mouse monoclonal antibody raised against amino acids 31-160 mapping near the N-terminus of VPAC1 of human origin.

# PRODUCT

Each vial contains 200  $\mu g$  IgG\_{2a} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

VPAC1 (B-4) is available conjugated to agarose (sc-377152 AC), 500 µg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-377152 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-377152 PE), fluorescein (sc-377152 FITC), Alexa Fluor<sup>®</sup> 488 (sc-377152 AF488), Alexa Fluor<sup>®</sup> 546 (sc-377152 AF546), Alexa Fluor<sup>®</sup> 594 (sc-377152 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-377152 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-377152 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-377152 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

# **STORAGE**

Store at 4° C, \*\*D0 NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

#### **APPLICATIONS**

VPAC1 (B-4) is recommended for detection of VPAC1 of human origin by Western Blotting (starting dilution 1:100, 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), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

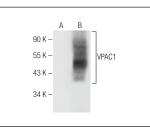
Suitable for use as control antibody for VPAC1 siRNA (h): sc-40281, VPAC1 shRNA Plasmid (h): sc-40281-SH and VPAC1 shRNA (h) Lentiviral Particles: sc-40281-V.

Molecular Weight of degylcosylated VPAC1: 47 kDa.

Molecular Weight of gylcosylated VPAC1: 58 kDa.

Positive Controls: TE671 cell lysate: sc-2416, Caki-1 cell lysate: sc-2224 or VPAC1 (h): 293T Lysate: sc-116969.

# DATA





VPAC1 (B-4): sc-377152. Western blot analysis of VPAC1 expression in non-transfected: sc-117752 (**A**) and human VPAC1 transfected: sc-116969 (**B**) 293T whole cell lysates. VPAC1 (B-4): sc-377152. Immunoperoxidase staining of formalin fixed, paraffin-embedded human skeletal muscle tissue showing cytoplasmic staining of myocytes.

# **SELECT PRODUCT CITATIONS**

- 1. Ulkumen, B., et al. 2022. Role of VPAC1 and VPAC2 receptors in the etiology of pregnancy rhinitis: an experimental study in rats. Braz. J. Otorhinolaryngol. 88: 505-510.
- Kitayama, E., et al. 2023. Functional expression of IP, 5-HT<sub>4</sub>, D<sub>1</sub>, A<sub>2A</sub>, and VIP receptors in human odontoblast cell line. Biomolecules 13: 879.

#### **RESEARCH USE**

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

#### **PROTOCOLS**

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