



AVP Receptor V3 (m): 293 Lysate: sc-178317

BACKGROUND

Vasopressin (AVP), the antidiuretic hormone, is a cyclic nonpeptide that is involved in the regulation of body fluid osmolality. AVP mediates its effects through a family of G protein-coupled receptors, the vasopressin receptors type V1a, V2 and V3 (also designated V1b). The AVP receptor V1a is responsible for several functions, including blood vessel constriction, liver glycogenolysis and platelet adhesion. It is detected as a full length protein and a shorter protein, which results from proteolytic cleavage of its amino terminus. The V1a receptor is coupled to $G_{q/11}$ protein, which increases the intracellular calcium concentration. The human AVP receptor V2 gene maps to chromosome Xq28 and is expressed in lung and kidney. Mutations in the V2 receptor result in nephrogenic diabetes insipidus (NDI), a rare X-linked disorder characterized by the inability of the kidney to concentrate urine in response to AVP. The AVP Receptor V2 activates the G_s protein and the cyclic AMP second messenger system. The AVP receptor V3 is preferentially expressed in the pituitary and stimulates the release of adrenocorticotrophic hormone (ACTH) in response to AVP by mobilizing intracellular calcium stores. AVP receptor antagonists may have potential therapeutic effects in hypertension, congestive heart failure, nephrotic syndrome and ACTH-secreting tumors.

REFERENCES

1. Thibonnier, M., Auzan, C., Madhun, Z., Wilkins, P., Berti-Mattera, L. and Clauser, E. 1994. Molecular cloning, sequencing, and functional expression of a cDNA encoding the human V1a vasopressin receptor. *J. Biol. Chem.* 269: 3304-3310.
2. Sugimoto, T., Saito, M., Mochizuki, S., Watanabe, Y., Hashimoto, S. and Kawashima, H. 1994. Molecular cloning and functional expression of a cDNA encoding the human V1b vasopressin receptor. *J. Biol. Chem.* 269: 27088-27092.
3. Fay, M.J., Du, J., Yu, X. and North, W.G. 1996. Evidence for expression of vasopressin V2 receptor mRNA in human lung. *Peptides* 17: 477-481.
4. Phalipou, S., Cotte, N., Carnazzi, E., Seyer, R., Mahe, E., Jard, S., Barberis, C. and Mouillac, B. 1997. Mapping peptide-binding domains of the human V1a vasopressin receptor with a photoactivatable linear peptide antagonist. *J. Biol. Chem.* 272: 26536-26544.
5. Mircic, G.M., Beleslin, D.B. and Jankovic, S.M. 1998. Hormones of the posterior region of the hypophyseal gland. *Srp. Arh. Celok. Lek.* 126: 111-118.
6. Birnbaumer, M. 1999. Vasopressin receptor mutations and nephrogenic diabetes insipidus. *Arch. Med. Res.* 30: 465-474.
7. Thibonnier, M., Coles, P., Thibonnier, A. and Shoham, M. 2001. The basic and clinical pharmacology of nonpeptide vasopressin receptor antagonists. *Annu. Rev. Pharmacol. Toxicol.* 41: 175-202.
8. Morello, J.P. and Bichet, D.G. 2001. Nephrogenic diabetes insipidus. *Annu. Rev. Physiol.* 63: 607-630.

STORAGE

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

CHROMOSOMAL LOCATION

Genetic locus: Avpr1b (mouse) mapping to 1 E4.

PRODUCT

AVP Receptor V3 (m): 293 Lysate represents a lysate of mouse AVP Receptor V3 transfected 293 cells and is provided as 100 µg protein in 200 µl SDS-PAGE buffer.

APPLICATIONS

AVP Receptor V3 (m): 293 Lysate is suitable as a Western Blotting positive control for mouse reactive AVP Receptor V3 antibodies. Recommended use: 10-20 µl per lane.

Control 293 Lysate: sc-110760 is available as a Western Blotting negative control lysate derived from non-transfected 293 cells.

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.