AT₁ (F-3): sc-365493



The Power to Question

BACKGROUND

Angiotensin II (Ang II) is an important physiological effector of blood pressure and volume regulation through vasoconstriction, aldosterone release, sodium uptake and thirst stimulation. Although Ang II interacts with two types of cell surface receptors, AT₁ and AT₂, most of the major cardiovascular effects seem to be mediated through AT₁. Molecular cloning of the AT₁ protein has shown it to be a member of the G protein-associated seven transmembrane protein receptor family. Ang II treatment of cells results in activation of several signal transduction pathways as evidenced by tyrosine phosphorylation of several proteins and induction of others. PLC γ is phosphorylated after 30 seconds of treatment with Angiotensin II, indicating this as an early signal transduction event. Ang II treatment also stimulates phosphorylation of Shc, FAK and MAP kinases and induces MKP-1, indicating stimulation of growth factor pathways. Ang II stimulation through AT₁ has been shown to activate the JAK/Stat pathway involving a direct interaction between JAK2 and AT₁ as demonstrated by coimmunprecipitation. The AT₁ receptor has no cytoplasmic kinase domain, but is able to function as a substrate for Src kinases and has several putative phosphorylation sites.

REFERENCES

- Murphy, T.J., et al. 1991. Isolation of a cDNA encoding the vascular type-1 Angiotensin II receptor. Nature 351: 233-236.
- 2. Tsuda, T., et al. 1991. Vasoconstrictor-induced protein-tyrosine phosphorylation in cultured vascular smooth muscle cells. FEBS Lett. 285: 44-48.
- Duff, J.L., et al. 1993. Angiotensin II induces 3CH134, a protein-tyrosine phosphatase, in vascular smooth muscle cells. J. Biol. Chem. 268: 26037-26040.
- Timmermans, P.B., et al. 1993. Angiotensin II receptors and Angiotensin II receptor antagonists. Pharmacol. Rev. 45: 205-251.
- Marrero, M.B., et al. 1994. Angiotensin II stimulates tyrosine phosphorylation of phospholipase C-γ 1 in vascular smooth muscle cells. J. Biol. Chem. 269: 10935-10939.

CHROMOSOMAL LOCATION

Genetic locus: AGTR1 (human) mapping to 3q24; Agtr1b (mouse) mapping to 3 A2.

SOURCE

 ${\rm AT_1}$ (F-3) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 10-29 within an N-terminal extracellular domain of ${\rm AT_1}$ of human origin.

PRODUCT

Each vial contains 200 μg lgM in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

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

APPLICATIONS

AT $_1$ (F-3) is recommended for detection of AT $_1$ of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], 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).

AT₁ (F-3) is also recommended for detection of AT₁ in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for AT_1 siRNA (h): sc-29750, AT_1 siRNA (m): sc-29751, AT_1 siRNA (r): sc-155992, AT_1 shRNA Plasmid (h): sc-29750-SH, AT_1 shRNA Plasmid (m): sc-29751-SH, AT_1 shRNA Plasmid (r): sc-155992-SH, AT_1 shRNA (h) Lentiviral Particles: sc-29750-V, AT_1 shRNA (m) Lentiviral Particles: sc-29751-V and AT_1 shRNA (r) Lentiviral Particles: sc-155992-V.

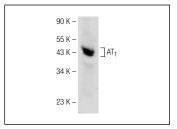
Molecular Weight of AT₁: 43 kDa.

Positive Controls: CCRF-CEM cell lysate: sc-2225, THP-1 cell lysate: sc-2238 or A549 cell lysate: sc-2413.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-mouse IgM-HRP: sc-2064 (dilution range: 1:500-1:5,000), TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein L PLUS-Agarose: sc-2336 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use goat anti-mouse IgM-FITC: sc-2082 (dilution range: 1:100-1:400) or goat anti-mouse IgM-TR: sc-2983 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



 ${\rm AT_1}$ (F-3): sc-365493. Western blot analysis of ${\rm AT_1}$ expression in A549 whole cell lysate.

SELECT PRODUCT CITATIONS

 Karpe, P.A., et al. 2012. Insulin resistance induces a segmental difference in thoracic and abdominal aorta: differential expression of AT₁ and AT₂ receptors. J. Hypertens. 30: 132-146.

RESEARCH USE

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

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