p-DARPP-32 (Thr 75): sc-135688



The Power to Question

BACKGROUND

Dopaminergic signaling pathways, which are essential for multiple brain functions, are abnormal in several neurological disorders, such as schizophrenia, Parkinson's disease and drug abuse. DARPP-32 is abundant in neurons that receive dopaminergic input. Activation of PKA and the consequent phosphorylation of DARPP-32 on Thr 34 occurs in response to dopamine acting upon D1-like receptors. Dopamine interaction with D2-like receptors results in the inhibition of PKA activation, the activation of protein phosphatase 2B and the consequent dephosphorylation of DARPP-32 at Thr 34. Phosphorylated DARPP-32 at Thr 34 is a potent inhibitor of PP-1. Phosphorylation of DARPP-32 on Ser 137 by casein kinase inhibits the dephosphorylation of Thr 34 by calcineurin. Phosphorylation of DARPP-32 on Thr 75 by CdK5 inhibits PKA by a competitive mechanism *in vitro*. Decreasing the phosphorylation of DARPP-32 Thr 75 increases the dopamine-induced phosphorylation of PKA substrates.

REFERENCES

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- Desdouits, F., et al. 1995. Dopamine- and cAMP-regulated phosphoprotein DARPP-32: phosphorylation of Ser 137 by casein kinase I inhibits dephosphorylation of Thr 34 by calcineurin. Proc. Natl. Acad. Sci. USA 92: 2682-2685.
- Nishi, A., et al. 1997. Bidirectional regulation of DARPP-32 phosphorylation by dopamine. J. Neurosci. 17: 8147-8155.
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CHROMOSOMAL LOCATION

Genetic locus: PPP1R1B (human) mapping to 17q12; Ppp1r1b (mouse) mapping to 11 D.

SOURCE

p-DARPP-32 (Thr 75) is a rabbit polyclonal antibody raised against a short amino acid sequence containing phosphorylated Thr 75 of DARPP-32 of rat origin.

PRODUCT

Each vial contains IgG in 100 μ l of 10 mM HEPES with 150 mM NaCl, 50% glycerol and < 0.1% BSA.

APPLICATIONS

p-DARPP-32 (Thr 75) is recommended for detection of Thr 75 phosphorylated DARPP-32 of mouse, rat and human origin by Western Blotting (starting dilution to be determined by researcher, dilution range 1:100-1:5000) and immunoprecipitation [1-2 μ l per 100-500 μ g of total protein (1 ml of cell lysate)].

Suitable for use as control antibody for DARPP-32 siRNA (h): sc-35173, DARPP-32 siRNA (m): sc-35174, DARPP-32 shRNA Plasmid (h): sc-35173-SH, DARPP-32 shRNA Plasmid (m): sc-35174-SH, DARPP-32 shRNA (h) Lentiviral Particles: sc-35173-V and DARPP-32 shRNA (m) Lentiviral Particles: sc-35174-V.

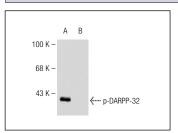
Molecular Weight of p-DARPP-32: 32 kDa.

Positive Controls: rat caudate nucleus tissue extract.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit lgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit lgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto B Blocking Reagent: sc-2335 (use 50 mM NaF, sc-24988, as diluent), Western Blotting Luminol Reagent: sc-2048 and Lambda Phosphatase: sc-200312A. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).

DATA



p-DARPP-32 (Thr 75): sc-135688. Western blot analysis of DARPP-32 phosphorylation in untreated (**A**) and lambda protein phosphatase-treated (**B**) rat caudate nucleus tissue extracts.

STORAGE

For immediate and continuous use, store at 4° C for up to one month. For sporadic use, freeze in working aliquots in order to avoid repeated freeze/thaw cycles. If turbidity is evident upon prolonged storage, clarify solution by centrifugation.

RESEARCH USE

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

PROTOCOLS

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