SANTA CRUZ BIOTECHNOLOGY, INC.

JNK3 (m): 293T Lysate: sc-121159



BACKGROUND

c-Jun N-terminal kinases (JNKs) phosphorylate and augment transcriptional activity of c-Jun. JNKs originate from three genes that yield 10 isoforms through alternative mRNA splicing, including JNK1 α 1, JNK1 β 1, JNK2 α 1, JNK2 β 1 and JNK3 α 1, which represent the p46 isoforms, and JNK1 α 2, JNK1 β 2, JNK2 α 2, JNK2 β 2 and JNK3 β 2, which represent the p54 isoforms. JNKs coordinate cell responses to stress and influence regulation of cell growth and transformation. The human JNK1 (PRKM8, SAPK1, MAPK8) gene maps to chromosome 10q11.22 and shares 83% amino acid identity with JNK2. JNK1 is necessary for normal activation and differentiation of CD4 helper T (TH) cells into TH1 and TH2 effector cells. Capsaicin activates JNK1 and p38 in Ras-transformed human breast epithelial cells. Nitrogen oxides (NOx) upregulate JNK1 in addition to c-Fos, c-Jun and other signaling kinases, including MEKK1 and p38. JNK3 (MK10, MAPK10, PRKM10) is activated by pro-inflammatory cytokines and environmental stresss by phosphorylating transcription factors such as c-Jun and ATF-2. This is important for AP-1 transcriptional activity regulation. JNK3 is crucial for neuronal apoptosis (stressinduced).

REFERENCES

- Gupta, S., et al. 1996. Selective interaction of JNK protein kinase isoforms with transcription factors. EMBO J. 15: 2760-2770.
- Dong, C., et al. 1998. Defective T cell differentiation in the absence of JNK1. Science 282: 2092-2095.
- Xie, X., et al. 1998. Crystal structure of JNK3: a kinase implicated in neuronal apoptosis. Structure 6: 983-991.
- Potapova, O., et al. 2000. Inhibition of c-Jun N-terminal kinase 2 expression suppresses growth and induces apoptosis of human tumor cells in a p53-dependent manner. Mol. Cell. Biol. 20: 1713-1722.
- 5. Dong, C., et al. 2000. JNK is required for effector T cell function but not for T cell activation. Nature 405: 91-94.
- Lisnock, J., et al. 2000. Activation of JNK3α1 requires both MKK4 and MKK7: kinetic characterization of *in vitro* phosphorylated JNK3α1. Biochemistry 39: 3141-3148.
- Dreskin, S.C., et al. 2001. Isoforms of Jun kinase are differentially expressed and activated in human monocyte/macrophage (THP-1) cells. J. Immunol. 166: 5646-5653.
- Han, S.Y., et al. 2002. Differential gene regulation by specific gain-offunction JNK1 proteins expressed in Swiss 3T3 fibroblasts. J. Biol. Chem. 277: 47167-47174.
- 9. Chou, F.P., et al. 2002. Induced proliferation of human MRC-5 cells by nitrogen oxides via direct and indirect activation of MEKK1, JNK, and p38 signals. Toxicol. Appl. Pharmacol. 181: 203-208.

CHROMOSOMAL LOCATION

Genetic locus: Mapk10 (mouse) mapping to 5 E5.

RESEARCH USE

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

PRODUCT

JNK3 (m): 293T Lysate represents a lysate of mouse JNK3 transfected 293T cells and is provided as 100 μ g protein in 200 μ l SDS-PAGE buffer.

APPLICATIONS

JNK3 (m): 293T Lysate is suitable as a Western Blotting positive control for mouse reactive JNK3 antibodies. Recommended use: 10-20 μl per lane.

Control 293T Lysate: sc-117752 is available as a Western Blotting negative control lysate derived from non-transfected 293T cells.

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.

PROTOCOLS

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