9223

ATF-2 Control Cell Extracts

🗸 200 uL (Controls for 10 western blots)



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For Research Use Only. Not For Use In Diagnostic Procedures.

Product Includes	Product #	Quantity
ATF-2 Control Cell Extracts (3T3 untreated)	68481	200 ul
ATF-2 Control Cell Extracts (3T3 +anisomycin)	82288	200 ul

Storage: Supplied in SDS Sample Buffer: 62.5 mM Tris-HCI (pH 6.8 at 25°C), 2% w/v SDS, 10% glycerol, 50 mM DTT, 0.01% w/v phenol red or bromophenol blue. Store at -20°C or at -80°C for long term storage.

Please visit www.cellsignal.com for a complete listing of recommended complementary products.

Background: The transcription factor ATF-2 (also called CRE-BP1) binds to both AP-1 and CRE DNA response elements and is a member of the ATF/CREB family of leucine zipper proteins (1). ATF-2 interacts with a variety of viral oncoproteins and cellular tumor suppressors and is a target of the SAPK/JNK and p38 MAP kinase signaling pathways (2-4). Various forms of cellular stress, including genotoxic agents, inflammatory cytokines, and UV irradiation, stimulate the transcriptional activity of ATF-2. Cellular stress activates ATF-2 by phosphorylation of Thr69 and Thr71 (2-4). Both SAPK and p38 MAPK have been shown to phosphorylate ATF-2 at these sites in vitro and in cells transfected with ATF-2. Mutations of these sites result in the loss of stress-induced transcription by ATF-2 (2-4). In addition, mutations at these sites reduce the ability of E1A and Rb to stimulate gene expression via ATF-2 (2).

Description: Nonphosphorylated ATF-2 Control Cell Extracts: Total extracts from NIH/3T3 cells, to serve as a negative control. Supplied in SDS Sample Buffer.

Phosphorylated ATF-2 Control Cell Extracts: Total extracts from NIH/3T3 cells, treated with Anisomycin #2222 at 25 ug/ml for 30 minutes to serve as a positive control. Supplied in SDS Sample Buffer.

Directions for Use: Boil for 3 minutes prior to use. Load 20 µl of phosphorylated and nonphosphorylated ATF-2 Control Cell Extracts per lane.

Background References:

Applications Kev:

- (1) Abdel-Hafiz, H.A. et al. (1992) Mol. Endocrinol. 6, 2079-2089.
- (2) Gupta, S. et al. (1995) Science 267, 389-393.
- (3) van Dam, H. et al. (1995) EMBO J. 14, 1798-1811.
- (4) Livingstone, C. et al. (1995) EMBO J. 14, 1785-1797.



Western blot analysis of ATF-2 Control Cell Extracts using Phospho-ATF-2 (Thr69/71) Antibody #9225 (upper) and ATF-2 (20F1) Rabbit mAb #9226 (lower).

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W-Western IP—Immunoprecipitation IHC—Immunohistochemistry IC—Immunocytochemistry IF-Immunofluorescence F—Flow cytometry E—ELISA D-DELEIA® Mk-monkey Species Cross-Reactivity Key: H—human M—mouse R—rat Hm—hamster Mi—mink C—chicken X—Xenopus Z—zebra fish B-bovine All-all species expected Species enclosed in parentheses are predicted to react based on 100% sequence homology.

ATF-2 Signaling Pathway