Stat1 Control Cell Extracts

Controls for 10 western blots



Orders = 877-616-CELL (2355) orders@cellsignal.com Support = 877-678-TECH (8324) info@cellsignal.com Web = www.cellsignal.com

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Background: Stat1, while activated in response to a large number of ligands (1), appears to be essential for responsiveness to IFN- α and IFN- γ (2,3). Phosphorylation of Stat1 at Tyr701 induces Stat1 dimerization, nuclear translocation and DNA binding (4). Stat1 has two isoforms, Stat1 α (91 kDa) and the splice variant Stat1 β (84 kDa). In most cells, both isoforms are activated by IFN- α , but only Stat1 α is activated by IFN- γ . Stat1 has been found to be inappropriately activated in many tumors (5). In addition to tyrosine phosphorylation, Stat1 is phosphorylated through a p38 mitogen-activated protein kinase (MAPK)-dependent pathway at Ser727 in response to IFN- α and other cellular stresses (6). Serine phosphorylation may be required for the maximal induction of Stat1-mediated gene activation.

Description:

Nonphosphorylated Stat1 Control Cell Extracts: Total cell extracts from HeLa cells prepared without treatment serve as a negative control. Supplied in SDS Sample Buffer.

Phosphorylated Stat1 Control Cell Extracts: Total cell extracts from HeLa cells prepared with 100 ng/ml interferonalpha 5 minute treatment serve as a positive control. Supplied in SDS Sample Buffer. Western Blots: CST recommends using 10 ul of phosphorylated and nonphosphorylated Stat1 control cell extracts as controls. Boil for 3 minutes prior to the first use.

Background References:

(1) Heim, M.H. (1999) *J. Recept. Signal. Transduct. Res.* 19, 75–120.

(2) Durbin, J.E. et al. (1996) Cell 84, 443-450.

(3) Meraz, M.A. et al. (1996) Cell 84, 431-442.

(4) Ihle, J.N. et al. (1994) *Trends Biochem. Sci.* 19, 222–227.

(5) Frank, D.A. (1999) *Mol. Med.* 5, 432–456.
(6) Wen, Z. et al. (1995) *Cell* 82, 241–250.

Entrez-Gene ID #6772 UniProt ID #P42224

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 bromophenol blue. Store at -20°C. *Store at -80°C long term.*

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