Technical Data Sheet Purified Mouse Anti-Stat1

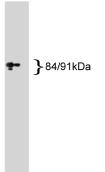
Product Information	
Material Number:	610116
Size:	150 µg
Concentration:	250 μg/ml
Clone:	1/Stat1
Immunogen:	Human Stat1 aa. 1-194
Isotype:	Mouse IgG1
Reactivity: Target MW:	QC Testing: Human Tested in Development: Chicken, Dog, Frog, Mouse, Rat 91/84 kDa
Storage Buffer:	Aqueous buffered solution containing BSA, glycerol, and $\leq 0.09\%$ sodium azide.

Description

Stat (Signal transducer and activators of transcription) proteins are critical mediators of the biologic activity of cytokines, including interleukins, interferons, erythropoietin, and growth factors. Ligand-receptor interaction leads to activation of constitutively associated JAK family kinases and subsequent recruitment/activation of Stat proteins by tyrosine phosphorylation. Active Stat proteins then move to the nucleus to promote transcription of cytokine-inducible genes. Seven Stat proteins have been cloned, each of which is differentially expressed and/or activated in a cytokine-specific and cell type-specific manner. Stat1 and Stat2 are components of the ISGF3 (Interferon-Stimulated Gene Factor 3) complex, which is the primary transcription activator induced by the binding of the interferon to a specific cell-surface receptor. Stat1 has two alternatively spliced isoforms, 91-kDa Stat1 α and 84 kDa Stat1 β ; Stat1 α has 38 additional C-terminal amino acids. In response to the binding of IFN α , IFN γ , EGF, PDGF, or CSF-1 to their respective receptors, the Stat1 subunits become tyrosine-phosphorylated at Y701, and the complex is translocated to the nucleus. This results in the formation of an active complex that includes the DNA-binding p48 subunit. This complex is responsible for modulating the transcription of the interferon-stimulated genes (ISGs). Thus, phosphorylation of Y701 in Stat1 occurs in response to growth factors and cytokines, and is essential for normal transcriptional activity of the ISGF3 complex.

The 1/Stat1 monoclonal antibody recognizes the N-terminus of human Stat1 (both isoforms), regardless of phosphorylation status.

This antibody is routinely tested by western blot analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.





Immunofluorescent staining of human fibroblast cells.

123

Western blot analysis of Stat1 on a A431 cell lysate. Lane 1: 1:250, lane 2: 1:500, lane 3: 1:1000 dilution of the anti- Stat1 antibody.

BD Biosciences

bdbiosciences.c	om				
United States 877.232.8995	Canada 888.259.0187 cific contact infor	Europe 32.53.720.550	Japan 0120.8555.90 osciences.com/how	Asia Pacific 65.6861.0633	Latin America/Caribbear 55.11.5185.9995
of any patents. BD I use of our products	Biosciences will not b s. Purchase does not i nponent of another p	e held responsible for nclude or carry any rig roduct. Any use of th	trued as a recommend patent infringement of ght to resell or transfer is product other than t ictly prohibited.	or other violations that this product either as	may occur with the a stand-alone

BD, BD Logo and all other trademarks are the property of Becton, Dickinson and Company. ©2006 BD



Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at -20° C.

Application Notes

Application

Western blot	Routinely Tested
Immunofluorescence	Tested During Development
Immunohistochemistry	Tested During Development
Immunoprecipitation	Tested During Development

Suggested Companion Products

Catalog Number	Name	Size	Clone
611447	A431 Cell Lysate	500 μg	(none)
554002	HRP Goat Anti-Mouse Igs	1.0 ml	(none)
554001	FITC Goat Anti-Mouse Igs	0.5 mg	Polyclonal

Product Notices

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- 2. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

References

Doan TN, Ali MS, Bernstein KE. Tyrosine kinase activation by the angiotensin II receptor in the absence of calcium signaling. J Biol Chem. 2001;

276(24):20954-20958.(Clone-specific: Immunoprecipitation, Western blot)

Dumoutier L, Louahed J, Renauld JC. Cloning and characterization of IL-10-related T cell-derived inducible factor (IL-TIF), a novel cytokine structurally related to IL-10 and inducible by IL-9. J Immunol. 2000; 164(4):1814-1819.(Clone-specific: Gel shift)

Dupuis S, Dargemont C, Fieschi C, et al. Impairment of mycobacterial but not viral immunity by a germline human STAT1 mutation. Science. 2001;

293(5528):300-303.(Clone-specific: Immunohistochemistry, Immunoprecipitation, Western blot)

Fu XY, Schindler C, Improta T, Aebersold R, Darnell JE Jr. The proteins of ISGF-3, the interferon alpha-induced transcriptional activator, define a gene family involved in signal transduction. *Proc Natl Acad Sci U S A*. 1992; 89(16):7840-7843.(Biology)

Sadowski HB, Shuai K, Darnell JE Jr, Gilman MZ. A common nuclear signal transduction pathway activated by growth factor and cytokine receptors. *Science*. 1993; 261(5129):1739-1744.(Biology)