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STAT1 Polyclonal Antibody

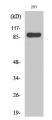
catalog number: E-AB-32977

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description	
Reactivity	Human;Mouse;Rat
Immunogen	Synthesized peptide derived from human Stat1 around the non-phosphorylation site
	of Ser727.
Host	Rabbit
Isotype	IgG
Purification	Affinity purification
Buffer	Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer, 0.5% protein
	protectant and 50% glycerol.
Applications	Recommended Dilution

WB	1:500-1:2000	
IHC	1:100-1:300	
IF	1:50-1:200	

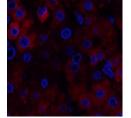
Data





Western Blot analysis of 293 cells using Stat1 Polyclonal Antibody at dilution of 1:2000.

Observed-MW:87 kDa Calculated-MW:87 kDa



Immunohistochemistry of paraffin-embedded Human liver cancer tissue using Stat1 Polyclonal Antibody at dilution of 1:200.

Immunofluorescence analysis of Human liver tissue using Stat1 Polyclonal Antibody at dilution of 1:200.

Preparation & Storage	
Storage	Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.
Shipping	The product is shipped with ice pack, upon receipt, store it immediately at the
	temperature recommended.

Background

For Research Use Only

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Signal transducer and activator of transcription that mediates signaling by interferons (IFNs). Following type I IFN (IFNalpha and IFN-beta) binding to cell surface receptors, Jak kinases (TYK2 and JAK1) are activated, leading to tyrosine phosphorylation of STAT1 and STAT2. The phosphorylated STATs dimerize, associate with ISGF3G/IRF-9 to form a complex termed ISGF3 transcription factor, that enters the nucleus. ISGF3 binds to the IFN stimulated response element (ISRE) to activate the transcription of interferon stimulated genes, which drive the cell in an antiviral state. In response to type II IFN (IFN-gamma), STAT1 is tyrosine- and serine-phosphorylated. It then forms a homodimer termed IFN-gammaactivated factor (GAF), migrates into the nucleus and binds to the IFN gamma activated sequence (GAS) to drive the expression of the target genes, inducing a cellular antiviral state.

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