

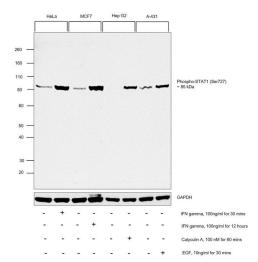


# Phospho-STAT1 (Ser727) Recombinant Rabbit Monoclonal Antibody (2H10)

<b>Product Details</b>	
Size	100 μL
Species Reactivity	Human
Host/Isotype	Rabbit / IgG
Expression system	HEK293 cells
Class	Recombinant Monoclonal
Туре	Antibody
Clone	2H10
Conjugate	Unconjugated
Immunogen	A synthesized peptide derived from human Phospho-STAT1 (S727)
Form	Liquid
Concentration	0.64 mg/mL
Purification	Affinity chromatography
Storage buffer	PBS, pH 7.4, with 50% glycerol
Contains	0.02% sodium azide
Storage conditions	-20°C or -80°C if preferred
RRID	AB_2812014

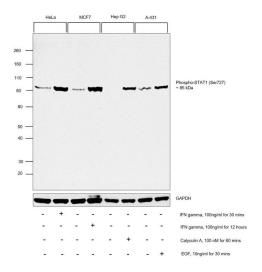
Applications	Tested Dilution	Publications
Western Blot (WB)	1:500-1:5,000	-
Immunohistochemistry (Paraffin) (IHC (P))	1:50-1:200	-
Immunocytochemistry (ICC/IF)	1:20-1:200	-
ELISA (ELISA)	Assay-Dependent	-

# Product Images For Phospho-STAT1 (Ser727) Recombinant Rabbit Monoclonal Antibody (2H10)



### Phospho-STAT1 (Ser727) Antibody (MA5-33198)

Altered expression of proteins upon cell treatment demonstrates antibody specificity. Western blot using Phospho-STAT1 (Ser727) Recombinant Rabbit Monoclonal Antibody (Product # MA5-33198), shows increased expression of proteins in HeLa and MCF7 on IFN gamma treatment, and A-431 upon EGF treatment, and induction of proteins in Hep G2 on Calyculin A treatment. {TM}



# Phospho-STAT1 (Ser727) Antibody (MA5-33198) in WB

Western blot was performed using Anti-Phospho-STAT1 (Ser727) Recombinant Rabbit Monoclonal Antibody (Product # MA5-33198) and a 85 kDa band corresponding to Signal transducer and activator of transcription 1-alpha/beta was observed in HeLa, MCF7 and A-431 cells which increased with IFN gamma and EGF treatments, and was induced in Hep G2 upon Calyculin A treatment. Modified whole cell extracts (1% SDS) (30 µg lysate) of HeLa (Lane 1), HeLa treated with IFN gamma (100 ng/mL for 30 minutes) (Lane 2), MCF7 (Lane 3), MCF7 treated with IFN gamma (100 ng/mL for 12 hours) (Lane 4), Hep G2 (Lane 5), Hep G2 treated with Calvculin A (100 nm for 60 minutes) (Lane 6), A-431 (Lane 7) and A-431 treated with EGF (10 ng/mL for 30 minutes) (Lane 8) were electrophoresed using NuPAGE™ 4-12% Bis-Tris Protein Gel (Product # NP0322BOX). Resolved proteins were then transferred onto a Nitrocellulose membrane (Product # IB23002) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with the primary antibody (1:1000 dilution) and detected by chemiluminescence with Goat anti-Rabbit IgG (Heavy Chain) Superclonal™ Recombinant Secondary Antibody, HRP (Product # A27036, 1: 4000 dilution) using the iBright FL 1000 (Product # A32752). Chemiluminescent detection was performed using Novex® ECL Reagent Kit (Product # WP20005).

# Phospho-STAT1(Ser727) HeLa (IFN gamma treated) D C Composite Phospho-STAT1(Ser727) HeLa (Untreated) Phospho-STAT1(Ser727) HeLa (Untreated) F-Actin F-Actin F-Actin F-Actin F-Actin F-Actin F-Actin

## Phospho-STAT1 (Ser727) Antibody (MA5-33198) in ICC/IF

Immunofluorescence analysis of Signal transducer and activator of transcription 1-alpha/beta was performed using 70% confluent log phase HeLa and HeLa cells treated with IFN gamma (100 ng/mL for 30 minutes). The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 15 minutes, and blocked with 2% BSA for 45 minutes at room temperature. The cells were labeled with Phospho-STAT1 (Ser727) Recombinant Rabbit Monoclonal Antibody (Product # MA5-33198) at 1:100 dilution in 0.1% BSA, incubated at 4 degree celsius overnight and then labeled with Donkey anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488 (Product # A32790), (1:2000 dilution), for 45 minutes at room temperature (Panel a: Green). Nuclei (Panel b:Blue) were stained with ProLong™ Diamond Antifade Mountant with DAPI (Product # P36962). F-actin (Panel c: Red) was stained with Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing Nuclear localization. Panel e represents untreated HeLa cells with no signal. Panel f represents control cells with no primary antibody to assess background. The images were captured at 60X magnification.

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