

Stat1(Phospho Tyr701) Polyclonal Antibody

Description

Product type Primary Antibody

Code BT-AP01644

Host Rabbit

Isotype IgG

Size 20ul, 50ul, 100ul

Immunogen The antiserum was produced against synthesized peptide derived from human STAT1 around the

phosphorylation site of Tyr701. AA range:668-717

Mol wt 87335

Species reactivity Human, Mouse, Rat, Monkey

Clonality Polyclonal

Recommended application IF, ICC, WB, IHC-p, IP, ELISA

Concentration 1 mg/ml

Full name Signal transducer and activator of transcription 1-alpha/beta

Synonyms Signal transducer and activator of transcription 1-alpha/beta; STAT1; Signal transducer and activator of

transcription 1-alpha/beta; Transcription factor ISGF-3 components p91/p84

This product is for research use only, not for use in human, therapeutic or diagnostic procedure.

Background

The protein encoded by this gene is a member of the STAT protein family. In response to cytokines and growth factors STAT family members are phosphorylated by the receptor associated kinases and then form homo- or heterodimers that translocate to the cell nucleus where they act as transcription activators. This protein can be activated by various ligands including interferon-alpha interferon-gamma EGF PDGF and IL6. This protein mediates the expression of a variety of genes which is thought to be important for cell viability in response to different cell stimuli and pathogens. Two alternatively spliced transcript variants encoding distinct isoforms have been described.

Recommended Dilution

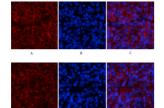
WB: 1: 500 - 1: 2000

IP: 2 - 5 ug: mg

IHC-p: 1: 100 - 1: 300 IF: 1: 50 - 1: 200 ELISA: 1: 10000

Not yet tested in other applications.

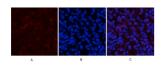
Images

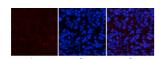


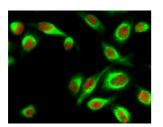
Immunofluorescence analysis of Hela cell. Stat1 (phospho Tyr701) Polyclonal Antibody(Red) was diluted at 1:200(4°C overnight). HER2 Monoclonal Antibody(11H9)(Green) was diluted at 1:200(4°C overnight).

Immunofluorescence analysis of rat-spleen tissue. 1,Stat1 (phospho Tyr701) Polyclonal Antibody(Red) was diluted at 1:200(4°C overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI.

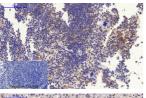
Picture C: merge of A+B

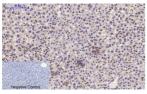


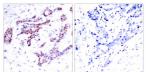


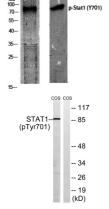














Immunofluorescence analysis of rat-spleen tissue. 1,Stat1 (phospho Tyr701) Polyclonal Antibody(Red) was diluted at 1:200(4°C overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

Immunofluorescence analysis of mouse-spleen tissue. 1,Stat1 (phospho Tyr701) Polyclonal Antibody(Red) was diluted at 1:200(4°C overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

Immunofluorescence analysis of mouse-spleen tissue. 1,Stat1 (phospho Tyr701) Polyclonal Antibody(Red) was diluted at 1:200(4°C overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

Immunohistochemical analysis of paraffin-embedded Rat-lung tissue. 1,Stat1 (phospho Tyr701) Polyclonal Antibody was diluted at 1:200(4°C overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.

Immunohistochemical analysis of paraffin-embedded Mouse-liver tissue. 1,Caspase-1 Polyclonal Antibody was diluted at 1:200(4°C overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.

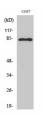
Immunohistochemical analysis of paraffin-embedded Mouse-spleen tissue. 1,Stat1 (phospho Tyr701) Polyclonal Antibody was diluted at 1:200(4°C overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.

Immunohistochemistry analysis of paraffin-embedded human breast carcinoma, using STAT1 (Phospho-Tyr701) Antibody. The picture on the right is blocked with the phospho peptide.

Western blot analysis of lysates from 1) MOUSE-BRAIN, 2) SY-SY5X ,3) COS7 cells, (Green) primary antibody was diluted at 1:1000, 4°C overnight, secondary antibody was diluted at 1:10000, 37°C 1hour. (Red) Actin β Monoclonal Antibody(5B7) was diluted at 1:5000 as loading control, 4°C overnight, secondary antibody was diluted at 1:10000, 37°C 1hour.

Western Blot analysis of various cells using Phospho-Stat1 (Y701) Polyclonal Antibody diluted at 1:1000

Western Blot analysis of COS7 cells using Phospho-Stat1 (Y701) Polyclonal Antibody diluted at 1:1000



Western blot analysis of lysates from COS7 cells, using STAT1 (Phospho-Tyr701) Antibody. The lane on the right is blocked with the phospho peptide.

Storage

-20°C for 1 year

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