

Mouse Monoclonal Antibody to

**STAT1 (phospho-Ser 727)**

**clone 12C5**

**Order No.:** 0176-100/STAT1-12C5  
**Size (µg)** 100  
**Lot No.:** 0176S



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02/160307F

Isotype	Species Reactivity	Applications	Mol. Weight	Ref.Cell Line	Epitope	Immunogen
IgG1	human	WB, ELISA	92 kDa	HepG2	phosphoserine 727 L P M pS P E E	phosphopeptide conjugated to KLH

**Background and Specificity:**

The STAT proteins serve as both cytoplasmic signal transducers and nuclear activators of transcription. STATs are mediators involved in cytokine signalling. In response to a specific cytokine signal, STAT proteins are phosphorylated on conserved tyrosine residues. Phosphorylated STAT proteins dimerize via their SH2 domains and move to the nucleus. The STAT dimers bind to specific DNA elements resulting in transcriptional regulation of downstream target genes.

STAT1 is activated by phosphorylation at serine 727. The phosphorylation state of Ser 727 regulates transcription and apoptosis. STAT1 can bind to DNA as heterodimer with STAT3.

**Mab STAT1-12C5** specifically recognizes STAT1 phosphorylated at Ser 727. The antibody does not crossreact with the non-phosphorylated form of STAT1 nor with unrelated serine-phosphorylated proteins. Mab STAT1-12C5 is suitable for Western blot and ELISA applications.

**Related Products**

- mab to STAT3 (phospho-Tyr 705)**  
#0036-100/STAT3-9E12
- mab to STAT3 (phospho-Ser 727)**  
#0145-100/STAT3-23G5
- mab to STAT5 A/B (phospho-Tyr 695/699)**  
#0121-100/STAT5-5G4
- mab to STAT6 (phosph-Tyr 641)**  
#0079-100/STAT6-16E12
- mab to STAT6 (aa 630-650)**  
#0063-100/STAT6-8C12

**Purification:** The antibody was purified from serum-free cell culture supernatant by subsequent thiophilic adsorption and size exclusion chromatography.

**Formulation:** liquid; 0.1mg/ml in in PBS/0.09% Na-Azide/PEG and Sucrose/50% Glycerol

**Reconstitution:**

**Stability:** Aliquote and store at -20°C up to 1 year.

**Avoid repeated freeze / thaw cycles.**

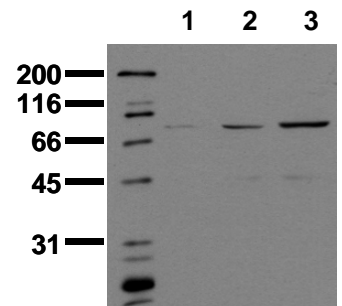
**Positive Control:** #0813: Cell lysate from EGF-treated HepG2 cells

**Immunoblotting:** 0.5 µg/ml for HRPO/ECL detection  
**Recommended blocking buffer:** Casein/Tween 20 based blocking and blot incubation buffer, e.g. nanoTools product #3031-500/CPPT or #3031-3000/CPPT.

**Immunoprecipitation:** ND

**Immunocytochemistry:** ND

**ELISA:** use at 0.05 µg/ml



**Phosphospecificity**

Whole cell extracts of control (1), EGF stimulated (2) or pervanadate treated (3) A549 tumor cells were applied to SDS-PAGE (ca 20.000 cells per lane) and transferred to a PVDF membrane. The immunoblot was probed with mab STAT1-12C5 (0.5 µg/ ml) for 1h at RT and developed by ECL (exp. time: 30 sec).

**All products are supplied for research and investigational use only. Not for use in humans or laboratory animals.**