

Mouse Monoclonal Antibody to

STAT1 (phospho-Ser 727)

clone 12C5

Size (µg) Lot No.: 100

0176S





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Lot No.	.:	0176S			02/160307F			
Isotype	Species Reactivity	Applications	Mol. Weight	Ref.Cell Line	Epitope		Immunogen	
lgG1	human	WB, ELISA	92 kDa	HepG2	phosphoserir L P M pS P E		phosphopeptide conjugated to KLH	
Background and Specificity:						Related Pro	oducts	
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.						#0036-100/STAT3 mab to STAT3 #0145-100/STAT3 mab to STAT5 695/699) #0121-100/STAT5 mab to STAT0 #0079-100/STAT6	3 (phospho-Ser 727) -23G5 5 A/B (phospho-Tyr -5G4 6 (phosph-Tyr 641) -16E12 6 (aa 630-650)	
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					1 2 3	
Immunoblo	Re blo		king buffer: Ca	sein/Tween 20 bas g. nanoTools prodi ⊇T.		200 — 116 — 66 — 45 — 31 —		
Immunopre	ecipitation: NE)				Phosphospecifici	ty	
-	tochemistry: ND)				Whole cell extracts	of control (1), EGF stimulated (2) or	
ELISA:	-	e at 0.05 µg/ml				SDS-PAGE (ca 20 PVDF membrane. STAT1-12C5 (0.5)	d (3) A549 tumor cells were applied to .000 cells per lane) and transferred to a The immunoblot was probed with mab µg/ ml) for 1h at RT and developed by	
	All products	are supplied for	esearch and in	vestigational		ECL (exp. time: 30	sec).	

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