Mouse Monoclonal Antibody to

STAT1 (phospho-Ser 727)

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clone 12C5								
Order No.:		0176-100/STAT1-12C5				ŏ		
Size (μ	a)	100				ğ		
Lot No.:		0176S				8		
					56	02/1603		
Isotype	Species Reactivity	Applications	Mol. Weight	Ref.Cell Line	Epitope	707	Immunogen	
lgG1	human	WB, ELISA	92 kDa	HepG2	phosphoserii	ne /2/	phosphopeptide conjugated to KLH	
					L P M pS P E	ΞE		
Destaura						Polotod Pro	duata	
Backgroun	nd and Specificity:					Related Pro	Daucts	
	proteins serve as both						3 (phospho-Tyr 705)	
<u>transcription</u> . STATs are mediators involved in cytokine signalling. In response to a specific cytokine signal, STAT proteins are phosphorylated on conserved tyrosine residues.				ecific	#0036-100/STAT3-9E12 mab to STAT3 (phospho-Ser 727)			
Phosphoryl	ated STAT proteins d rs bind to specific DN	merize via their SI	H2 domains and	move to the nucle		#0145-100/STAT3-23G5 mab to STAT5 A/B (phospho-Tyr		
target gene	s.			C C		695/699)	o Alb (phospho-ry)	
	ctivated by phosphory anscription and apopt					#0121-100/STAT5-5G4		
0						mab to STAT6 (phosph-Tyr 641) #0079-100/STAT6-16E12		
	1-12C5 specifically re- act with the non-phos				body does	mab to STAT #0063-100/STAT6	6 (aa 630-650) -8012	
serine-phos	sphorylated proteins. I				ISA	#0005-100/51A10	-0012	
applications	5.							
su		The antibody was purified from serum-free cell culture supernatant by subsequent thiophilic adsorption and size exclusion chromatography.						
		iquid; 0.1mg/ml in in PBS/0.09% Na-Azide/PEG and						
		Sucrose/50% Glycerol						
Reconstitu								
Stability: A		Aliquote and store at -20°C up to 1 year.						
Av		Avoid repeated freeze / thaw cycles.						
Positive Co	ontrol: #08	13: Cell lysate from	m EGF-treated H	HepG2 cells			1 2 3	
Immunoble		µg/ml for HRPO/E				200-	-	
	<u>Re</u>	commended bloc	king buffer: Ca	sein/Tween 20 bas g. nanoTools produ	sed	116—	=	
	#30	31-500/CPPT or #	#3031-3000/CPF	PT.		66—	-	
						45 —	-	
						31 —		
						31 —	=	
Immunopr	ecipitation: ND					31 — Phosphospecifici	-	
-	ecipitation: ND tochemistry: ND					Phosphospecific	• ity s of control (1), EGF stimulated (2) or dd (3) A549 tumor cells were applied to	

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PVDF membrane. The immunoblot was probed with mab STAT1-12C5 (0.5 μ g/ ml) for 1h at RT and developed by ECL (exp. time: 30 sec).