Phospho-Glycogen Synthase (Ser641) Antibody Store at -20C **Cell Signaling** TECHNOLOGY® Orders: 877-616-CELL (2355) orders@cellsignal.com Support: 877-678-TECH (8324) . 68 80 Web: info@cellsignal.com cellsignal.com 3 Trask Lane | Danvers | Massachusetts | 01923 | USA

## For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: WB, IP	Reactivity: H M R	Sensitivity: Endogenous	<b>MW (kDa):</b> 85 to 90	Source: Rabbit	UniProt ID: #P13807	Entrez-Gene Id: 2997
Product Usage Information	Ар	Application Dilution				
mormation	We	estern Blotting			1:1000	
	Imr	nunoprecipitation			1:100	
Storage		plied in 10 mM sodi C. Do not aliquot the		5), 150 mM NaCl, 10	00 μg/ml BSA and 50% g	Jycerol. Store at –
Specificity / Sensit	isofo	Phospho-Glycogen Synthase (Ser641) Antibody detects endogenous levels of both muscle and liver isoforms of glycogen synthase only when phosphorylated at serine 640 or 641 of the muscle and liver isoforms, respectively.				
Species predicted react based on 100 sequence homolog	0%	se				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser641 of human liver glycogen synthase. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Glycogen is a polysaccharide of glucose and serves as an energy storage in mammalian muscle and liver (1). Glycogen synthase catalyzes the rate-limiting step of glycogen biosynthesis and has two major isoforms in mammals: muscle isoform (glycogen synthase 1, GYS1) and liver isoform (glycogen synthase 2, GYS2), respectively (1). Glycogen synthase kinase- $3\alpha$ (GSK- $3\alpha$ ) and glycogen synthase kinase- $3\beta$ (GSK- $3\beta$ ) phosphorylate glycogen synthase at multiple sites in its C-terminus (Ser641, Ser645, Ser649, and Ser653), inhibiting its activity (2,3). Hypoxia alters glycogen metabolism including temporal changes of GYS1 expression and phosphorylation in cancer cells, suggesting the role of metabolic reprogramming of glycogen metabolism in cancer growth (1).				
Background Refer	2. M	avaro, E. et al. (201 lora, A. et al. (2005) ensen, J. et al. (2012	FEBS Lett 579, 36	32-8.	, E82-9.	
Species Reactivity	Spec	ies reactivity is dete	ermined by testing i	n at least one appro	oved application (e.g., we	estern blot).
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X 0.1% Tween $\$ 20 at 4°C with gentle shaking, overnight.				% w/v BSA, 1X TBS,
Applications Key	WB:	WB: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity K	<b>X</b> : Xe		B: bovine Dg: dog	Pg: pig Sc: S. cere	s <b>Mi:</b> mink <b>C:</b> chicken Dr evisiae <b>Ce:</b> C. elegans <b>H</b>	-
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Limited Uses						

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