## 59851

## Glycogen Synthase 1 (GYS1) Antibody



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

<b>Applications:</b> W, IP	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 84	Source/Isotype: Rabbit	UniProt ID: #P13807	Entrez-Gene Id: 2997
Product Usage Information	•	<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Glycogen Synthase 1 (GYS1) Antibody detects endogenous levels of total muscle glycogen synthase protein.				
Species predicted to react based on 100% sequence homology		Horse				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the sequence of human 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 $\alpha$ (GSK-3 $\alpha$ ) 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 References		1. Favaro, E. et al. (2012) <i>Cell Metab</i> 16, 751-64. 2. Mora, A. et al. (2005) <i>FEBS Lett</i> 579, 3632-8. 3. Jensen, J. et al. (2012) <i>Am J Physiol Endocrinol Metab</i> 303, E82-9.				
Species Reacti	vity	Species reactivity is det	termined by testin	g in at least one approve	ed application (e.g.,	western blot).
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity Key		H: Human M: Mouse R: Rat Mk: Monkey				
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