

# Phospho-HNF1 $\alpha$ (Ser247) Antibody



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

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W	H	Endogenous	81	Rabbit	#P20823	6927

Product Usage Information	Application	Dilution
	Western Blotting	1:1000
<b>Storage</b>	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.	
<b>Specificity/Sensitivity</b>	HNF1 $\alpha$ (Ser247) Antibody recognizes endogenous levels of HNF1 $\alpha$ protein only when phosphorylated at Ser247.	
<b>Species predicted to react based on 100% sequence homology</b>	Mouse, Rat	
<b>Source / Purification</b>	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser247 of human HNF1 $\alpha$ protein. Antibodies are purified by protein A and peptide affinity chromatography.	
<b>Background</b>	Hepatocyte nuclear factor 1 $\alpha$ (HNF1 $\alpha$ , also known as TCF1 or MODY3) is a transcription factor that plays a role in the tissue-specific regulation of liver gene expression (1). Research has shown that heterogeneous mutations of HNF1 $\alpha$ are linked to maturity onset diabetes of the young (MODY) (2). Recent studies indicate that increased concentrations of free fatty acids can reduce the expression of FoxA2/HNF3 $\beta$ and HNF1 $\alpha$ in pancreatic $\beta$ -cells and lead to their nuclear exclusion, resulting in symptoms of several metabolic syndromes (3). HNF1 $\alpha$ is inhibited through Akt2-mediated phosphorylation at Ser247 and is a negative regulator of PPAR $\gamma$ gene transcription. Research studies have shown that inhibition of PPAR $\gamma$ is beneficial in steatosis-associated liver cancer in mouse models (4).	
<b>Background References</b>	<ol style="list-style-type: none"> <li>1. Yamagata, K. et al. (1996) <i>Nature</i> 384, 455-8.</li> <li>2. Lehto, M. et al. (1997) <i>J Clin Invest</i> 99, 582-91.</li> <li>3. Ohtsubo, K. et al. (2011) <i>Nat Med</i> 17, 1067-75.</li> <li>4. Patitucci, C. et al. (2017) <i>J Clin Invest</i> 127, 1873-1888.</li> </ol>	

<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).
<b>Western Blot Buffer</b>	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween@ 20 at 4°C with gentle shaking, overnight.
<b>Applications Key</b>	<b>W:</b> Western Blotting
<b>Cross-Reactivity Key</b>	<b>H:</b> Human
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