Phospho-IRS-1 (Tyr1222) Antibody



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Applications: React	,	MW (kDa): 180	Source/Isotype: Rabbit	UniProt ID: #P35568	Entrez-Gene Id 3667	
Product Usage Information	Application Western Blotting			Dilution 1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity	Phospho-IRS-1 (Tyr1222) Antibody detects endogenous levels of IRS-1 only when phosphorylat Tyr1222. The antibody may cross-react with other activated receptor tyrosine kinases (RTKs) are docking proteins.					
Species predicted to re based on 100% sequer homology						
Source / Purification	corresponding to re	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr1222 of human IRS-1. Antibodies are purified by protein A and peptide affinity chromatography.				
Background	1 contains multiple to containing proteins also contains over 3 phosphorylated by J family (7). The PKC a respectively (8,9). Ph	cyrosine phosphoryla that mediate the me 0 potential serine/thi NK (5) and IKK (6) wh and mTOR pathways i osphorylation of IRS	of the major substrates tion motifs that serve as cabolic and growth-pron econine phosphorylatior ile Ser789 is phosphoryl nediate phosphorylatio 1 at Ser1101 is mediate 1 a potential mechanism	s docking sites for S noting functions of n sites. Ser307 of IR lated by SIK-2, a me n of IRS-1 at Ser612 d by PKC0 and resu	H2-domain insulin (2-4). IRS-1 S-1 is mber of the AMPK and Ser636/639, ilts in an inhibition	
	Tyr1222 provides a d	Phosphorylation of tyrosine 1222 of IRS-1 was identified in insulin stimulated cells (11). Phosphorylate Tyr1222 provides a docking site for the SH2 domain of PTP2C, which may mediate dephosphorylation of IRS-1 and lead to negative feedback of insulin signaling (12).				
Background Reference	2. Sun, X.J. et al. (1993). Myers Jr., M.G. et al. (Wang, L.M. et al. (1997). G. Gao, Z. et al. (2002). Horike, N. et al. (2002). Horike, N. et al. (2004). De Fea, K. and Ruito. Li, Y. et al. (2004). Sun, X.J. et al. (19	1. Sun, X.J. et al. (1991) <i>Nature</i> 352, 73-77. 2. Sun, X.J. et al. (1992) <i>J. Biol. Chem.</i> 267, 22662-22672. 3. Myers Jr., M.G. et al. (1993) <i>Endocrinology</i> 132, 1421-1430. 4. Wang, L.M. et al. (1993) <i>Science</i> 261, 1591-1594. 5. Rui, L. et al. (1997) <i>J. Clin. Invest.</i> 107, 181-189. 6. Gao, Z. et al. (2002) <i>J. Biol. Chem.</i> 277, 48115-48121. 7. Horike, N. et al. (2003) <i>J. Biol. Chem.</i> 278, 18440-18447. 8. Ozes, O.N. et al. (2001) <i>Proc. Natl. Acad. Sci. USA</i> 98, 4640-4645. 9. De Fea, K. and Ruth, R.A. (1997) <i>Biochemistry</i> 36, 12939-12947. 10. Li, Y. et al. (2004) <i>J. Biol. Chem.</i> 279, 45304-45307. 11. Sun, X.J. et al. (1993) <i>Mol. Cell. Biol.</i> 13, 7418-7428. 12. Rocchi, S. et al. (1995) <i>Endocrinology</i> 136, 5291-5297.				

Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer 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.

Applications Key W: Western Blotting

Cross-Reactivity Key H: Human

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