## AS160 Antibody Image: Display the product of the p

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Applications: W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 160	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #O60343	Entrez-Gene Id: 9882
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 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		AS160 Antibody detects endogenous levels of total AS160 protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Cys1286 of human AS160. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Insulin is a major hormone controlling critical energy functions, such as glucose and lipid metabolism. Insulin binds to and activates the insulin receptor (IR) tyrosine kinase, which phosphorylates and recruits adaptor proteins. The signaling pathway initiated by insulin and its receptor stimulates glucose uptake in muscle cells and adipocytes through translocation of the Glut4 glucose transporter from the cytoplasm to the plasma membrane (1). A 160 kDa substrate of the Akt Ser/Thr kinase (AS160, TBC1D4) is a Rab GTPase-activating protein that regulates insulin-stimulated Glut4 trafficking. AS160 is expressed in many tissues including brain, kidney, liver, and brown and white fat (2). Multiple Akt phosphorylation sites have been identified on AS160 <i>in vivo</i> , with five sites (Ser318, Ser570, Ser588, Thr642, and Thr751) showing increased phosphorylation following insulin treatment (2,3). Studies using recombinant AS160 demonstrate that insulin-stimulated phosphorylation of AS160 is a crucial step in Glut4 translocation (3) and is reduced in some patients with type 2 diabetes (4). The interaction of 14-3-3 regulatory proteins with AS160 phosphorylated at Thr642 is a necessary step for Glut4 translocation (5). Phosphorylation of AS160 by AMPK is involved in the regulation of contraction- stimulated Glut4 translocation (6).				
Background References		1. Watson, R.T. and Pessin, J.E. (2006) <i>Trends Biochem. Sci.</i> 31, 215-22. 2. Kane, S. et al. (2002) <i>J. Biol. Chem.</i> 277, 22115-8. 3. Sano, H. et al. (2003) <i>J. Biol. Chem.</i> 278, 14599-602. 4. Karlsson, H.K. et al. (2005) <i>Diabetes</i> 54, 1692-7. 5. Ramm, G. et al. (2006) <i>J. Biol. Chem.</i> 281, 29174-80. 6. Kramer, H.F. et al. (2006) <i>J. Biol. Chem.</i> 281, 31478-85.				
Species Reactiv	/ity	Species reactivity is det	ermined 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 BSA, 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				
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