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Atg12 Antibody



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Applications:Reactivity:W, IP, IF-ICM	Sensitivity: Endogenous	MW (kDa): 16, 55	Source/Isotype: Rabbit	UniProt ID: #Q9CQY1	Entrez-Gene Id: 67526	
Product Usage Information	Application Western Blotting Immunoprecipitation Immunofluorescence		istry)		Dilution 1:1000 1:50 1:100	
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	Atg12 Antibody detects endogenous levels of total free and Atg5 bound Atg12 proteins.					
Source / Purification	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of mouse Atg12. Antibodies are purified by peptide affinity chromatography.					
Background	Autophagy is a catabolic process for the autophagosomic-lysosomal degradation of bulk cytoplasmic contents (1,2). Autophagy is generally activated by conditions of nutrient deprivation but has also been associated with a number of physiological processes including development, differentiation, neurodegeneration, infection, and cancer (3). The molecular machinery of autophagy was largely discovered in yeast and referred to as autophagy-related (<i>Atg</i>) genes. Formation of the autophagosome involves a ubiquitin-like conjugation system in which Atg12 is covalently bound to Atg5 and targeted to autophagosome vesicles (4-6). This conjugation reaction is mediated by the ubiquitin E1-like enzyme Atg7 and the E2-like enzyme Atg10 (7,8).					
Background References	 Reggiori, F. and Klionsky, D.J. (2002) Eukaryot Cell 1, 11-21. Codogno, P. and Meijer, A.J. (2005) Cell Death Differ 12 Suppl 2, 1509-18. Levine, B. and Yuan, J. (2005) J Clin Invest 115, 2679-88. Mizushima, N. et al. (1998) J Biol Chem 273, 33889-92. Mizushima, N. et al. (1998) Nature 395, 395-8. Suzuki, K. et al. (2001) EMBO J 20, 5971-81. Tanida, I. et al. (1999) Mol Biol Cell 10, 1367-79. Shintani, T. et al. (1999) EMBO J 18, 5234-41. 					
Species Reactivity	Species reactivity is d	etermined by testing	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 IF-IC: Immunofluorescence (Immunocytochemistry)					
Cross-Reactivity Key	M: Mouse					
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