Atg12 (D88H11) Rabbit mAb

100 μl (10 western blots)



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Applications	Species Cross-Reactivity*	Molecular Wt.	Isotype	
W, IP	H, M, R, Mk	16, 55 kDa	Rabbit IgG**	

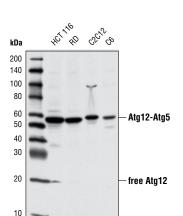
Background: Autophagy is a catabolic process for the autophagosomic-lysosomal degradation of bulk cytoplasmic contents (1,2). Autophagy is generally activated by conditions of nutrient deprivation but has also been associated with a number of physiological processes including development, differentiation, neurodegeneration, infection and cancer (3). The molecular machinery of autophagy was largely discovered in yeast and referred to as autophagy-related (Atg) genes. Formation of the autophagosome involves an 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).

Specificity/Sensitivity: Atg12 (D88H11) Rabbit mAb detects endogenous levels of total free and Atg5 bound Atg12 protein.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser36 of human Atg12 protein.

Background References:

- (1) Reggiori, F. and Klionsky, D.J. (2002) Eukaryot Cell 1,
- (2) Codogno, P. and Meijer, A.J. (2005) Cell Death Differ 12 Suppl 2, 1509-18,
- (3) Levine, B. and Yuan, J. (2005) J Clin Invest 115, 2679-
- (4) Mizushima, N. et al. (1998) J Biol Chem 273, 33889-
- (5) Mizushima, N. et al. (1998) Nature 395, 395-8.
- (6) Suzuki, K. et al. (2001) EMBO J 20, 5971-81.
- (7) Tanida, I. et al. (1999) Mol Biol Cell 10, 1367-79.
- (8) Shintani, T. et al. (1999) EMBO J 18, 5234-41.



Western blot analysis of extracts from various cell lines using Atg12 (D88H11) Rabbit mAb.

Entrez-Gene ID #9140 Swiss-Prot Acc. #094817

Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

*Species cross-reactivity is determined by western blot.

**Anti-rabbit secondary antibodies must be used to detect this antibody.

Recommended Antibody Dilutions:

Western blotting 1:1000 Immunoprecipitation 1:100

For application specific protocols please see the web page for this product at www.cellsignal.com.

Please visit www.cellsignal.com for a complete listing of recommended companion products.

IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v BSA, 1X TBS, 0.1% Tween-20 at 4°C with gentle shaking, overnight.