LC3A (D50G8) XP® Rabbit mAb



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Applications: W, IHC-P	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 14, 16	Source/Isotype: Rabbit IgG	UniProt ID: #Q9H492	Entrez-Gene Id 84557
Product Usage Information		Application Western Blotting Immunohistochemistry (Paraffin)		Dilution 1:1000 1:6400		
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.				
Specificity/Sensitivity		LC3A (D50G8) $XP^{\$}$ Rabbit mAb detects endogenous levels of total LC3A protein. This antibody may also react with LC3B.				
Species predicted to react based on 100% sequence homology		Monkey, Dog				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human LC3A.				
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 it has also been associated with a number of physiological processes including development, differentiation, neurodegenerative diseases, infection, and cancer (3). Autophagy marker Light Chain 3 (LC3) was originally identified as a subunit of microtubule-associated proteins 1A and 1B (termed MAP1LC3) (4) and subsequently found to contain similarity to the yeast protein Apg8/Aut7/Cvt5 critical for autophagy (5). Three human LC3 isoforms (LC3A, LC3B, and LC3C) undergo posttranslational modifications during autophagy (6-9). Cleavage of LC3 at the carboxy terminus immediately following synthesis yields the cytosolic LC3-I form. During autophagy, LC3-I is converted to LC3-II through lipidation by a ubiquitin-like system involving Atg7 and Atg3 that allows for LC3 to become associated with autophagic vesicles (6-10). The presence of LC3 in autophagosomes and the conversion of LC3 to the lower migrating form, LC3-II, have been used as indicators of autophagy (11).				
Background References		2. Codogno, P. and M 3. Levine, B. and Yuar 4. Mann, S.S. and Har 5. Lang, T. et al. (1998 6. Kabeya, Y. et al. (20 7. He, H. et al. (2003) 8. Tanida, I. et al. (20	eijer, A.J. (2005) <i>Cell</i> n, J. (2005) <i>J. Clin. In</i> v nmarback, J.A. (1994 d) <i>EMBO J.</i> 17, 3597-(100) <i>EMBO J.</i> 19, 572 <i>J. Biol. Chem.</i> 278, 2 d) <i>J. Biol. Chem.</i> 279 Biochem. Biophys. R (2000) <i>Nature</i> 408, 4	9, 5720-28. 278, 29278-87. n. 279, 47704-10. hys. Res. Commun. 339, 437-42. 408, 488-92.		
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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 IHC-P: Immunohistochemistry (Paraffin)

Cross-Reactivity Key

H: Human M: Mouse R: Rat

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