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## LC3A/B (D3U4C) XP<sup>®</sup> Rabbit mAb (Alexa Fluor<sup>®</sup> 594 Conjugate)

For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> IF-IC	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q9H492, #Q9GZQ8	<b>Entrez-Gene Id:</b> 84557, 81631
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<b>Product Usage Information</b>	<b>Application</b> Immunofluorescence (Immunocytochemistry)	<b>Dilution</b> 1:50
<b>Storage</b>	Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the antibody. Protect from light. Do not freeze.	
<b>Specificity/Sensitivity</b>	LC3A/B (D3U4C) XP <sup>®</sup> Rabbit mAb recognizes endogenous levels of total LC3A and LC3B proteins.	
<b>Species predicted to react based on 100% sequence homology</b>	Xenopus, Bovine, Dog, Pig	
<b>Source / Purification</b>	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Leu44 of human LC3B protein (conserved in LC3A).	
<b>Description</b>	This Cell Signaling Technology antibody is conjugated to Alexa Fluor <sup>®</sup> 594 fluorescent dye and tested in-house for direct immunofluorescent analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated LC3A/B (D3U4C) XP <sup>®</sup> Rabbit mAb #12741.	
<b>Background</b>	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).	
<b>Background References</b>	<ol style="list-style-type: none"> <li>1. Reggiori, F. and Klionsky, D.J. (2002) <i>Eukaryot. Cell</i> 1, 11-21.</li> <li>2. Codogno, P. and Meijer, A.J. (2005) <i>Cell Death Differ.</i> 12 Suppl 2, 1509-18.</li> <li>3. Levine, B. and Yuan, J. (2005) <i>J. Clin. Invest.</i> 115, 2679-88.</li> <li>4. Mann, S.S. and Hammarback, J.A. (1994) <i>J. Biol. Chem.</i> 269, 11492-97.</li> <li>5. Lang, T. et al. (1998) <i>EMBO J.</i> 17, 3597-607.</li> <li>6. Kabeya, Y. et al. (2000) <i>EMBO J.</i> 19, 5720-28.</li> <li>7. He, H. et al. (2003) <i>J. Biol. Chem.</i> 278, 29278-87.</li> <li>8. Tanida, I. et al. (2004) <i>J. Biol. Chem.</i> 279, 47704-10.</li> <li>9. Wu, J. et al. (2006) <i>Biochem. Biophys. Res. Commun.</i> 339, 437-42.</li> <li>10. Ichimura, Y. et al. (2000) <i>Nature</i> 408, 488-92.</li> <li>11. Kabeya, Y. et al. (2004) <i>J. Cell Sci.</i> 117, 2805-12.</li> </ol>	
<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).	
<b>Applications Key</b>	<b>IF-IC:</b> Immunofluorescence (Immunocytochemistry)	
<b>Cross-Reactivity Key</b>	<b>H:</b> Human <b>M:</b> Mouse <b>R:</b> Rat	
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