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50G8) XP <sup>®</sup> Rabbit mAb	4599	20 µl	14, 16 kDa	Rabbit IgG
11) XP <sup>®</sup> Rabbit mAb	3868	20 µl	14, 16 kDa	Rabbit IgG
bit IgG, HRP-linked Antibody	7074	100 µl		Goat
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ription	The Autophagosome Marker accumulation of autophagos	<i>,</i>		2

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than Storage 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody. 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 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 Atq7 and the E2-like enzyme Atq10 (7,8). Autophagy marker Light Chain 3 (LC3) was originally identified as a subunit of microtubule-associated proteins 1A and 1B (termed MAP1LC3) (9) and subsequently found to contain similarity to the yeast protein Apg8/Aut7/Cvt5 that is critical for autophagy (10). Three human LC3 isoforms (LC3A, LC3B, and LC3C) undergo posttranslational modifications during autophagy (11-14). 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 (11-15). 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 (16). **Background References** 1. Reggiori, F. and Klionsky, D.J. (2002) Eukaryot Cell 1, 11-21. 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, 2679-88. 4. Mizushima, N. et al. (1998) / Biol Chem 273, 33889-92. 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. 9. Mann, S.S. and Hammarback, J.A. (1994) J Biol Chem 269, 11492-7. 10. Lang, T. et al. (1998) EMBO J 17, 3597-607. 11. Kabeya, Y. et al. (2000) *EMBO J* 19, 5720-8. 12. Wu, J. et al. (2006) Biochem Biophys Res Commun 339, 437-42. 13. He, H. et al. (2003) / Biol Chem 278, 29278-87. 14. Tanida, I. et al. (2004) / Biol Chem 279, 47704-10. 15. Ichimura, Y. et al. (2000) Nature 408, 488-92.

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16. Kabeya, Y. et al. (2004) J Cell Sci 117, 2805-12.

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LC3B (D11) XP <sup>®</sup> Rabbit mAb	3868	20 µl	14, 16 kDa	

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