

GABARAP (E1J4E) Rabbit mAb



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Applications: W, IF-IC	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 14, 16	Source/Isotype: Rabbit IgG	UniProt ID: #O95166	Entrez-Gene Id 11337
Product Usage Information		Application Western Blotting			Dilution 1:1000	
		Immunofluorescence (Immunocytochemistry)			1:200 - 1:400	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
		For a carrier free (BSA and azide free) version of this product see product #64055.				
Specificity/Sensitivity		GABARAP (E1J4E) Rabbit mAb recognizes endogenous levels of total GABARAP protein. This antibody does not cross-react with other GABARAP family members.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Arg40 of human GABARAP protein.				
Background		GABA _A receptor associated protein (GABARAP) is an Atg8 family protein with a key role in autophagy, which was originally discovered as a protein associated with the GABA _A receptor regulating receptor trafficking to the plasma membrane (1). Proteins in this family, including microtubule-associated protein light chain 3 (LC3) and GATE-16 (GABARAPL2), become incorporated into the autophagosomal membranes following autophagic stimuli such as starvation (2). Like the other family members, GABARAP is cleaved at its carboxyl terminus, which leads to conjugation by either of the phospholipids phosphatidylethanolamine or phosphatidylserine (3,4). This processing converts GABARAP from a type I to a type II membrane bound form involved in autophagosome biogenesis. Processing of GABARAP involves cleavage by Atg4 family members (5,6) followed by conjugation by the E1 and E2 like enzymes Atg7 and Atg3 (7,8). GABARAPL1/GEC1, a protein that is highly related to GABARAP, was identified as an estrogen inducible gene, and is also associated with autophagosomes (9-11).				
Background References		1. Wang, H. et al. (1999) <i>Nature</i> 397, 69-72. 2. Shpilka, T. et al. (2011) <i>Genome Biol</i> 12, 226. 3. Kabeya, Y. et al. (2004) <i>J Cell Sci</i> 117, 2805-12. 4. Sou, Y.S. et al. (2006) <i>J Biol Chem</i> 281, 3017-24. 5. Tanida, I. et al. (2004) <i>J Biol Chem</i> 279, 36268-76. 6. Hemelaar, J. et al. (2003) <i>J Biol Chem</i> 278, 51841-50. 7. Tanida, I. et al. (2001) <i>J Biol Chem</i> 276, 1701-6. 8. Tanida, I. et al. (2002) <i>J Biol Chem</i> 277, 13739-44. 9. Chakrama, F.Z. et al. (2010) <i>Autophagy</i> 6, 495-505. 10. Pellerin, I. et al. (1993) <i>Mol Cell Endocrinol</i> 90, R17-21. 11. Vernier-Magnin, S. et al. (2001) <i>Biochem Biophys Res Commun</i> 284, 118-25.				
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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 IF-IC: Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key

H: Human M: Mouse R: Rat

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