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Rab7 (D95F2) XP<sup>®</sup> Rabbit mAb (HRP Conjugate)

	eactivity: Sensitiv M R Mk Endoge		Bource/Isotype: Rabbit IgG	<b>UniProt ID:</b> #P51149	Entrez-Gene Id: 7879		
Product Usage	Application			Dilution			
Information	Western Blottir	ng		1:1000			
Storage		Supplied in 136 mM NaCl, 2.6 mM KCl, 12 mM sodium phosphate (pH 7.4) dibasic, 2 mg/ml BSA, and 50% glycerol. Store at –20°C. Do not aliquot the antibodies.					
Specificity / Sensitivit	<b>y</b> Rab7 (D95F2) >	Rab7 (D95F2) XP $^{\otimes}$ Rabbit mAb (HRP Conjugate) detects endogenous levels of total Rab7 protein.					
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Glu188 of human Rab7 protein.					
Product Description	peroxidase (HR	This Cell Signaling Technology antibody is conjugated to the carbohydrate groups of horseradish peroxidase (HRP) via its amine groups. The HRP conjugated antibody is expected to exhibit the same species cross-reactivity as the unconjugated Rab7 (D95F2) XP® Rabbit mAb #9367.					
MW (kDa)		23					
Background	in late endosom membrane traffi growth receptor vacuoles (4-6). 6-phosphate rec	Rab7 and Rab9 are members of the Ras superfamily of small Rab GTPases (1). Both proteins are located in late endosomes, but exert different functions. Rab7 associates with the RIPL effector protein to control membrane trafficking from early to late endosomes and to lysosomes (2,3). Rab7 also helps to regulate growth receptor endocytic trafficking and degradation (3,4), and maturation of phagosome and autophagic vacuoles (4-6). Rab9 interacts with its effector proteins p40 and TIP47 (7,8) to promote the MPR (mannose 6-phosphate receptor)-associated lysosomal enzyme transport between late endosomes and the trans Golgi network (9,10).					
Background Referenc	2. Feng, Y. et al. 3. Méresse, S. e 4. Ceresa, B.P. a 5. Jäger, S. et a 6. Méresse, S. e 7. Díaz, E. et al. 8. Barbero, P. et 9. Lombardi, D.	<ol> <li>Zerial, M. and McBride, H. (2001) Nat Rev Mol Cell Biol 2, 107-17.</li> <li>Feng, Y. et al. (1995) J Cell Biol 131, 1435-52.</li> <li>Méresse, S. et al. (1995) J Cell Sci 108 (Pt 11), 3349-58.</li> <li>Ceresa, B.P. and Bahr, S.J. (2006) J Biol Chem 281, 1099-106.</li> <li>Jäger, S. et al. (2004) J Cell Sci 117, 4837-48.</li> <li>Méresse, S. et al. (1999) EMBO J 18, 4394-403.</li> <li>Díaz, E. et al. (1997) J Cell Biol 138, 283-90.</li> <li>Barbero, P. et al. (2002) J Cell Biol 156, 511-8.</li> <li>Lombardi, D. et al. (1993) EMBO J 12, 677-82.</li> <li>Riederer, M.A. et al. (1994) J Cell Biol 125, 573-82.</li> </ol>					
Species Reactivity	Species reactivit	y is determined by test	ing in at least one approv	red application (e.g., w	estern 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	WB: Western Bl	WB: Western Blotting					
Cross-Reactivity Key	X: Xenopus Z: ze	H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected					
Trademarks and Patents		Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc. XP is a registered trademark of Cell Signaling Technology, Inc.					

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