β-Arrestin 1/2 (D24H9) Rabbit mAb





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Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 50	Source/Isotype: Rabbit IgG	UniProt ID: #P49407, #P32121	Entrez-Gene Id: 408, 409		
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:200			
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.				l and less than		
Specificity/Sensitivity		β -Arrestin 1/2 (D24H9) Rabbit mAb detects endogenous level of total beta-arrestin 1 and beta-arrestin 2 proteins.						
Source / Purification		Monoclonal antibody is produced by immunizing animals with a recombinant protein containing the carboxy-terminal region of human β -arrestin 1.						
Background		Arrestin proteins function as negative regulators of G protein-coupled receptor (GPCR) signaling. Cognate ligand binding stimulates GPCR phosphorylation, which is followed by binding of arrestin to the phosphorylated GPCR and the eventual internalization of the receptor and desensitization of GPCR signaling (1). Four distinct mammalian arrestin proteins are known. Arrestin 1 (also known as S- arrestin) and arrestin 4 (X-arrestin) are localized to retinal rods and cones, respectively. Arrestin 2 (also known as β -arrestin 1) and arrestin 3 (β -arrestin 2) are ubiquitously expressed and bind to most GPCRs (2). β -arrestins function as adaptor and scaffold proteins and play important roles in other processes, such as recruiting c-Src family proteins to GPCRs in Erk activation pathways (3,4). β -arrestins are also involved in some receptor tyrosine kinase signaling pathways (5-8). Additional evidence suggests that β -arrestins translocate to the nucleus and help regulate transcription by binding transcriptional cofactors (9,10).						
Background R	eferences	 Shenoy, S.K. and Lefkowitz, R.J. (2005) <i>Sci STKE</i> 2005, cm10. Lefkowitz, R.J. and Shenoy, S.K. (2005) <i>Science</i> 308, 512-7. Luttrell, L.M. et al. (1999) <i>Science</i> 283, 655-61. Luttrell, L.M. et al. (1999) <i>Curr Opin Cell Biol</i> 11, 177-83. Luttrell, L.M. and Lefkowitz, R.J. (2002) <i>J Cell Sci</i> 115, 455-65. Waters, C. et al. (2004) <i>Semin Cell Dev Biol</i> 15, 309-23. Lefkowitz, R.J. and Whalen, E.J. (2004) <i>Curr Opin Cell Biol</i> 16, 162-8. Waters, C.M. et al. (2005) <i>Cell Signal</i> 17, 263-77. Kang, J. et al. (2005) <i>Cell Sci</i> 120, 213-8. Ma, L. and Pei, G. (2007) <i>J Cell Biochem</i> 116, 767-77. 						
Species Reacti	vity	Species reactivity is det	ermined by testin	g in at least one appro	ved application (e.g., w	vestern blot).		
Western Blot E	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.				5% w/v BSA, 1X		
Applications K	ey	W: Western Blotting IP: Immunoprecipitation						
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat Mk: Monkey						
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