

YAP (D8H1X) XP[®] Rabbit mAb (Alexa Fluor[®] 647 Conjugate)



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Applications: IF-IC, FC-FP	Reactivity: H M R Hm Mk	Sensitivity: Endogenous	Source/Isotype: Rabbit IgG	UniProt ID: #P46937	Entrez-Gene Id: 10413
Product Usage Information		Application Immunofluorescence (Ir Flow Cytometry (Fixed/P			Dilution 1:50 - 1:100 1:50
Storage		Supplied in PBS (pH 7.2), antibody. Protect from li		zide and 2 mg/ml BS	A. Store at 4°C. Do not aliquot the
Specificity/Sensitivity		YAP (D8H1X) XP [®] Rabbit mAb (Alexa Fluor [®] 647 Conjugate) recognizes endogenous levels of total YAP protein.			
Species predicted to react based on 100% sequence homology		Bovine, Horse, Guinea P	g		
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the carboxy terminus of human YAP protein. The epitope corresponds to a region surrounding Pro435 of human YAP isoform 1. This sequence region is 100% conserved among all known isoforms of human YAP protein.			
Description		This Cell Signaling Technology antibody is conjugated to Alexa Fluor [®] 647 fluorescent dye and tested in-house for direct flow cytometric analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated YAP (D8H1X) XP [®] Rabbit mAb #14074.			
Background		YAP (Yes-associated protein, YAP65) was first identified based on its ability to associate with the SH3 domain of Yes. It also binds to other SH3 domain-containing proteins such as Nck, Crk, Src, and Abl (1). In addition to the SH3 binding motif, YAP contains a PDZ interaction motif, a coiled-coil domain, and WW domains (2-4). While initial studies of YAP all pointed towards a role in anchoring and targeting to specific subcellular compartments, subsequent studies showed that YAP is a transcriptional coactivator by virtue of its WW domain interacting with the PY motif (PPxY) of the transcription factor PEBP2 and other transcription factors (5). In its capacity as a transcriptional co-activator, YAP is now widely recognized as a central mediator of the Hippo Pathway, which plays a fundamental and widely conserved role in regulating tissue growth and organ size (6-8). Phosphorylation at multiple sites (e.g., Ser109, Ser127) by LATS kinases promotes YAP translocation from the nucleus to the cytoplasm, where it is sequestered through association with 14-3-3 proteins (7-9). These LATS-driven phosphorylation events serve to prime YAP for subsequent phosphorylation by CK18/ε in an adjacent phosphodegron, triggering proteasomal degradation of YAP (10).			
Background Refe	erences	1. Sudol, M. (1994) <i>Onco</i> 2. Mohler, P.J. et al. (1999) 3. Espanel, X. and Sudol, 4. Sudol, M. et al. (1995) 5. Yagi, R. et al. (1999) <i>El</i> 6. Dong, J. et al. (2007) <i>C</i> 7. Zhao, B. et al. (2010) <i>C</i> 8. Zhao, B. et al. (2012) <i>Ce</i> 10. Zhao, B. et al. (2012) <i>Ce</i> 10. Zhao, B. et al. (2010)	a) J Cell Biol 147, 879-90. M. (2001) J Biol Chem 270 FEBS Lett 369, 67-71. MBO J 18, 2551-62. Fell 130, 1120-33. Fenes Dev 24, 862-74. Fenes Dev 21, 2747-61. Fell 150, 780-91.	6, 14514-23.	

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Applications Key

IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key

H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey

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