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TCF1/TCF7 (C63D9) Rabbit mAb (Alexa Fluor[®] 647 Conjugate)



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: FC-FP	Reactivity: H M	Sensitivity: Endogenous	Source/Isotype: Rabbit IgG	UniProt ID: #P36402	Entrez-Gene Id: 6932		
Product Usage Information		Application Flow Cytometry (Fixed/Permeabilized)		Dilution 1:50			
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the antibody. Protect from light. Do not freeze.					
Specificity/Sensiti	vity	TCF1/TCF7 (C63D9) Rabbit mAb (Alexa Fluor [®] 647 Conjugate) detects endogenous levels of total TCF1/TCF7 protein. This antibody does not recognize the dominant negative isoforms of TCF1/TCF7 lacking the amino-terminal β-catenin binding domain and does not cross-react with LEF1.					
Source / Purificati	on	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to a region surrounding Pro95 of human TCF1/TCF7 protein.					
Description		This Cell Signaling Technology antibody is conjugated to Alexa Fluor [®] 647 fluorescent dye and tested in-house for direct flow cytometry analysis in human cells. The antibody is expected to exhibit the same species cross-reactivity as the unconjugated TCF1/TCF7 (C63D9) Rabbit mAb #2203.					
Background		LEF1 and TCF are members of the high mobility group (HMG) DNA-binding protein family of transcription factors that consists of the following: Lymphoid Enhancer Factor 1 (LEF1), T Cell Factor 1 (TCF1/TCF7), TCF3/TCF7L1, and TCF4/TCF7L2 (1). LEF1 and TCF1/TCF7 were originally identified as important factors that regulate early lymphoid development (2) and act downstream in Wnt signaling. LEF1 and TCF bind to Wnt response elements to provide docking sites for β -catenin, which translocates to the nucleus to promote the transcription of target genes upon activation of Wnt signaling (3). LEF1 and TCF are dynamically expressed during development and aberrant activation of the Wnt signaling pathway is involved in many types of cancers, including colon cancer (4,5).					
		TCF1/TCF7 has several isoforms due to alternative splicing and transcription from an alternative promoter. The isoforms generated by the alternative promoter do not contain the amino-terminal β-catenin binding domain and therefore may function in a dominant negative manner (6). TCF1/TCF7 displays dynamic expression both in the total amount and the type of isoforms expressed in T cells during development and differentiation (7).					
Background Refer	rences	 Waterman, M.L. (2004) <i>Cancer Metastasis Rev</i> 23, 41-52. Schilham, M.W. and Clevers, H. (1998) <i>Semin Immunol</i> 10, 127-32. Brantjes, H. et al. (2002) <i>Biol Chem</i> 383, 255-61. Reya, T. and Clevers, H. (2005) <i>Nature</i> 434, 843-50. Logan, C.Y. and Nusse, R. (2004) <i>Annu Rev Cell Dev Biol</i> 20, 781-810. Waterman, M.L. (2004) <i>Cancer Metastasis Rev</i> 23, 41-52. Willinger, T. et al. (2006) <i>J Immunol</i> 176, 1439-46. 					
Species Reactivity	,	Species reactivity is determined by testing in at least one approved application (e.g., western blot).					
Applications Key		FC-FP: Flow Cytometry (Fixed/Permeabilized)					
Cross-Reactivity K	ey	H: Human M: Mouse					
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