TCF4/TCF7L2 (C48H11) Rabbit mAb



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#						
For Research Use Only. Not for Use in Diagnostic Procedures.						
Applications: W, W-S, IP, ChIP, hIP-seq, C&R, C&T	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 58, 79	Source/Isotype: Rabbit IgG	UniProt ID: #Q9NQB0	Entrez-Gene I 6934
Product Usage Information		For optimal ChIP and ChIP-seq results, use 10 μl of antibody and 10 μg of chromatin (approximately 4 x 10 ⁶ cells) per IP. This antibody has been validated using SimpleChIP® Enzymatic Chromatin IP Kits.				
		The CUT&RUN dilution was determined using CUT&RUN Assay Kit #86652.				
		The CUT&Tag dilution was determined using CUT&Tag Assay Kit #77552.				
		Application			Dilution	
		Western Blotting			1:1000	
		Simple Western™			1:10 - 1:50	
		Immunoprecipitation			1:50	
		Chromatin IP			1:50	
		Chromatin IP-seq			1:50	
		CUT&RUN			1:50	
		CUT&Tag			1:50	
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.				
Specificity/Sensitivity		TCF4/TCF7L2 (C48H11) Rabbit mAb detects endogenous levels of total TCF4/TCF7L2 protein.				
Species predicted to react based on 100% sequence homology		Mouse, Chicken				
Source / Purific	ation	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Leu331 of human TCF4/TCF7L2.				
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.				

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).

TCF4/TCF7L2 is expressed widely during development. Gene targeting studies indicate that TCF4/TCF7L2 is required to maintain the crypt stem cells of the small intestine (6,7). TCF4/TCF7L2 has several splicing isoforms that are expressed differentially in tissues and during cancer progression (8,9). Studies also indicate that a variant of the TCF4/TCF7L2 gene confers an increased risk of type 2 diabetes (10).

Background References

- 1. Waterman, M.L. (2004) Cancer Metastasis Rev 23, 41-52.
- 2. Schilham, M.W. and Clevers, H. (1998) Semin Immunol 10, 127-32.
- 3. Brantjes, H. et al. (2002) *Biol Chem* 383, 255-61.
- 4. Reya, T. and Clevers, H. (2005) Nature 434, 843-50.
- 5. Logan, C.Y. and Nusse, R. (2004) Annu Rev Cell Dev Biol 20, 781-810.
- 6. Cho, E.A. and Dressler, G.R. (1998) Mech. Dev. 77, 9-18.
- 7. Korinek, V. et al. (1998) Nat. Genet. 19, 379-383.
- 8. Howng, S.L. et al. (2004) *Int. J. Oncol.* 25, 1685-1692.
- 9. Shiina, H. et al. (2003) Clin. Cancer Res. 9, 2121-2132.
- 10. Grant, S.F. et al. (2006) Nat. Genet. 38, 320-323.

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 W-S: Simple Western™ IP: Immunoprecipitation ChIP: Chromatin IP ChIP-seq:

Chromatin IP-seq C&R: CUT&RUN C&T: CUT&Tag

Cross-Reactivity Key H: Human

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