Fra2 (D2F1E) Rabbit mAb



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Applications: W, IP, IHC-P, ChIP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 35-45	Source/Isotype: Rabbit IgG	UniProt ID: #P15408	Entrez-Gene Id: 2355
Product Usage Information		For optimal ChIP 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.				
		Application Western Blotting Immunoprecipitation Immunohistochemist Chromatin IP			1: 1:	7500
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.				
		For a carrier free (BSA and azide free) version of this product see product #51520.				
Specificity/Sensitivity		Fra2 (D2F1E) Rabbit mAb recognizes endogenous levels of total Fra2 protein. This antibody does not cross-react with c-Fos or Fra1.				
Species predicted to react based on 100% sequence homology		Bovine, Horse				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val245 of human Fra2 protein.				
Background		The Fos family of nuclear oncogenes includes c-Fos, FosB, Fos-related antigen 1 (FRA1), and Fos-related antigen 2 (FRA2) (1). While most Fos proteins exist as a single isoform, the FosB protein exists as two isoforms: full-length FosB and a shorter form, FosB2 (Delta FosB), which lacks the carboxy-terminal 101 amino acids (1-3). The expression of Fos proteins is rapidly and transiently induced by a variety of extracellular stimuli, including growth factors, cytokines, neurotransmitters, polypeptide hormones, and stress. Fos proteins dimerize with Jun proteins (c-Jun, JunB, and JunD) to form Activator Protein-1 (AP-1), a transcription factor that binds to TRE/AP-1 elements and activates transcription. Fos and Jun proteins contain the leucine-zipper motif that mediates dimerization and an adjacent basic domain that binds to DNA. The various Fos/Jun heterodimers differ in their ability to transactivate AP-1 dependent genes. In addition to increased expression, phosphorylation of Fos proteins by Erk kinases in response to extracellular stimuli may further increase transcriptional activity (4-6). Phosphorylation of c-Fos at Ser32 and Thr232 by Erk5 increases protein stability and nuclear localization (5). Phosphorylation of FRA1 at Ser252 and Ser265 by Erk1/2 increases protein stability and leads to overexpression of FRA1 in cancer cells (6). Following growth factor stimulation, expression of FosB and c-Fos in quiescent fibroblasts is immediate, but very short-lived, with protein levels dissipating after several hours (7). FRA1 and FRA2 expression persists longer, and appreciable levels can be detected in asynchronously growing cells (8). Deregulated expression of c-Fos, FosB, or FRA2 can result in neoplastic cellular transformation; however, Delta FosB lacks the ability to transform cells (2,3).				
Background References		 Tulchinsky, E. (2000) Histol Histopathol 15, 921-8. Dobrazanski, P. et al. (1991) Mol Cell Biol 11, 5470-8. Nakabeppu, Y. and Nathans, D. (1991) Cell 64, 751-9. Rosenberger, S.F. et al. (1999) J Biol Chem 274, 1124-30. Sasaki, T. et al. (2006) Mol Cell 24, 63-75. Basbous, J. et al. (2007) Mol Cell Biol 27, 3936-50. Kovary, K. and Bravo, R. (1991) Mol Cell Biol 11, 2451-9. Kovary, K. and Bravo, R. (1992) Mol Cell Biol 12, 5015-23. 				

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 IP: Immunoprecipitation IHC-P: Immunohistochemistry (Paraffin) ChIP: Chromatin

IF

Cross-Reactivity Key H: Human

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