

Phospho-SMAD2 (Ser465/Ser467) (E8F3R) Rabbit mAb



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Applications: W, W-F, IP, IF-IC, FC- FP, ChIP	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 60	Source/Isotype: Rabbit IgG	UniProt ID: #Q15796	Entrez-Gene Id: 4087
Product Usage Information		For optimal ChIP results, use 10 μ l of antibody and 10 μ g of chromatin (approximately 4 × 10 ⁶ cells) per IP. This antibody has been validated using SimpleChIP [®] Enzymatic Chromatin IP Kits.				
		Application			Dilution	
		Western Blotting			1:10	00
		Fluorescent Western			1:1000	
		Immunoprecipitation	1		1:10)
		Immunofluorescence (Immunocytochemistry)			1:400 - 1:1600	
		Flow Cytometry (Fixe	d/Permeabilized)			0 - 1:1600
		Chromatin IP			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.				
		For a carrier free (BSA and azide free) version of this product see product #94610.				
Specificity/Sensitivity		Phospho-Smad2 (Ser465/467) (E8F3R) Rabbit mAb recognizes endogenous levels of Smad2 protein when phosphorylated at Ser465 and Ser467.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser465/467 of human Smad2 protein.				
Background		Members of the SMAD family of signal transduction molecules are components of a critical intracellular pathway that transmit TGF-β signals from the cell surface into the nucleus. Three distinct classes of SMADs have been defined: the receptor-regulated SMADs (R-SMADs), which include SMAD1, 2, 3, 5, and 9; the common-mediator SMAD (co-SMAD), SMAD4; and the antagonistic or inhibitory SMADs (I-SMADs), SMAD6 and 7 (1-5). Activated type I receptors associate with specific R-SMADs and phosphorylate them on a conserved carboxy-terminal SSXS motif. The phosphorylated R-SMADs dissociate from the receptor and form a heteromeric complex with SMAD4, initiating translocation of the heteromeric SMAD complex to the nucleus. Once in the nucleus, SMADs recruit a variety of DNA binding proteins that function to regulate transcriptional activity (6-8).				
Background References		 Heldin, C.H. et al. (1997) Nature 390, 465-71. Attisano, L. and Wrana, J.L. (1998) Curr Opin Cell Biol 10, 188-94. Derynck, R. et al. (1998) Cell 95, 737-40. Massagué, J. (1998) Annu Rev Biochem 67, 753-91. Whitman, M. (1998) Genes Dev 12, 2445-62. Wrana, J.L. (2000) Sci STKE 2000, re1. Attisano, L. and Wrana, J.L. (2002) Science 296, 1646-7. Moustakas, A. et al. (2001) J Cell Sci 114, 4359-69. 				
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-F:** Fluorescent Western **IP:** Immunoprecipitation **IF-IC:** Immunofluorescence (Immunocytochemistry) **FC-FP:** Flow Cytometry (Fixed/Permeabilized) **ChIP:** Chromatin IP

H: Human M: Mouse R: Rat

Cross-Reactivity Key

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