

Phospho-SMAD1 (Ser463/465)/ SMAD5 (Ser463/465)/ SMAD9 (Ser465/467) (D5B10) Rabbit mAb



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

Applications: W, IP, IF-IC, FC-FP, ChIP	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 60	Source/Isotype: Rabbit IgG	UniProt ID: #Q99717, #Q15797, #O15198	Entrez-Gene Id: 4090, 4086, 4093
Product Usage Information		For optimal ChIP results, use 10 µl of antibody and 10 µg of ch IP. This antibody has been validated using SimpleChIP [®] Enzym Application Western Blotting Immunoprecipitation Immunofluorescence (Immunocytochemistry) Flow Cytometry (Fixed/Permeabilized) Chromatin IP			romatin (approximately 4 x 10 ⁶ cells) per atic Chromatin IP Kits. Dilution 1:1000 1:50 1:400 - 1:1600 1:200 - 1:800 1:50	
Storage		Supplied in 10 mM so 0.02% sodium azide. S	dium HEPES (pH 7.5 Store at –20°C. Do n	δ), 150 mM NaCl, 100 μ ot aliquot the antibody	g/ml BSA, 50% glycero v.	and less than
		For a carrier free (BSA	A and azide free) ver	sion of this product se	e product #67959.	
Specificity/Sen	sitivity	Phospho-SMAD1 (Ser463/465)/ SMAD5 (Ser463/465)/ SMAD9 (Ser465/467) (D5B10) Rabbit mAb recognizes endogenous levels of SMAD1 and SMAD5 protein when phosphorylated at Ser463/465 and SMAD9 (SMAD8) protein when phosphorylated at Ser465/467.				
Species predict based on 100% homology	ed to react sequence	Monkey				
Source / Purific	ation	Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser463/465 of human SMAD1 and SMAD5 protein.				
Background		Bone morphogenetic wide range of critical differentiation, and a kinase receptors. Liga these receptors (3-5). terminal motif SSXS, a phosphorylated SMAI they regulate the tran phosphorylate residu Ser206 recruits Smurf at this site also promo	proteins (BMPs) con processes including poptosis (1,2). BMP and binding induces They subsequently as well as SMAD5 ar Ds dimerize with the scription of target of es in the linker region f1 to the linker region otes SMAD1 transcription	nstitute a large family of morphogenesis, cell-f receptors are member multimerization, auto phosphorylate SMAD1 of SMAD9 (SMAD8) at t e coactivating SMAD4 a genes (5). MAP kinases on of SMAD1, including on and leads to the deg iptional activity by recr	of signaling molecules ate determination, pro s of the TGF-β superfar phosphorylation, and a at Ser463 and Ser465 heir corresponding sit nd translocate to the r and CDKs 8 and 9 are g Ser206. Phosphorylat pradation of SMAD1 (6) uiting YAP to the linker	that regulate a liferation, mily of Ser/Thr activation of in the carboxy- es. These nucleus, where also reported to ion of SMAD1 at . Phosphorylation region (7).
Background Re	ferences	1. Hogan, B.L. (1996) <i>Genes Dev</i> 10, 1580-94. 2. Hoodless, P.A. et al. (1996) <i>Cell</i> 85, 489-500. 3. Klemm, J.D. et al. (1998) <i>Annu Rev Immunol</i> 16, 569-92. 4. Kretzschmar, M. et al. (1997) <i>Genes Dev</i> 11, 984-95. 5. Whitman, M. (1998) <i>Genes Dev</i> 12, 2445-62. 6. Sapkota, G. et al. (2007) <i>Mol Cell</i> 25, 441-54. 7. Alarcón, C. et al. (2009) <i>Cell</i> 139, 757-69.				
Species Reactiv	vity	Species reactivity is do	etermined by testin	g in at least one approv	ved application (e.g., w	estern blot).
Western Blot B	uffer	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 Ke	₽y	W: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized) ChIP: Chromatin IP				

Cross-Reactivity Key	H: Human M: Mouse R: Rat		
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