

#8828

## Phospho-SMAD2 (Ser465/467)/SMAD3 (Ser423/425) (D27F4) Rabbit mAb



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

Applications: W	Reactivity: H M R Mk	Sensitivity:	<b>MW (kDa):</b> 52, 60	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID:	Entrez-Gene Id: 4088, 4087
VV	H IVI K IVIK	Endogenous	52, 60	Rabbit 1gG	#P84022, #Q15796	4088, 4087
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Phospho-SMAD2 (Ser465/467)/SMAD3 (Ser423/425) (D27F4) Rabbit mAb recognizes endogenous levels of SMAD2 protein when phosphorylated at Ser465 and Ser467. This antibody also recognizes endogenous levels of SMAD3 protein when phosphorylated Ser422 only or at both Ser423 and Ser425.				
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		2. Attisano, L. and Wra 3. Derynck, R. et al. (19 4. Massagué, J. (1998) 5. Whitman, M. (1998) 6. Wrana, J.L. (2000) <i>Sc</i> 7. Attisano, L. and Wra	et al. (1997) <i>Nature</i> 390, 465-71. and Wrana, J.L. (1998) <i>Curr Opin Cell Biol</i> 10, 188-94. et al. (1998) <i>Cell</i> 95, 737-40. J. (1998) <i>Annu Rev Biochem</i> 67, 753-91. 1. (1998) <i>Genes Dev</i> 12, 2445-62. 2000) <i>Sci STKE</i> 2000, re1. and Wrana, J.L. (2002) <i>Science</i> 296, 1646-7. A. et al. (2001) <i>J Cell Sci</i> 114, 4359-69.			
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**Species Reactivity** 

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

**Western Blot Buffer** 

 $IMPORTANT: For western \ blots, incubate \ membrane \ with \ diluted \ primary \ antibody \ in \ 5\% \ w/v \ nonfat$ 

dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** 

W: Western Blotting

**Cross-Reactivity Key** 

H: Human M: Mouse R: Rat Mk: Monkey

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