

**Phospho-Smad3 (Ser423/425)/Smad1
(Ser463/465) Antibody**

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Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
Endogenous	60	Rabbit	#Q15797, #P84022	4086, 4088

Storage	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at -20°C. Do not aliquot the antibody.
Specificity/Sensitivity	Phospho-Smad3 (Ser423/425)/Smad 1 (Ser463/465) Antibody detects endogenous levels of Smad3 only when phosphorylated at serines 423 and 425. The antibody cross-reacts with Smad1 when phosphorylated at serines 463 and 465 and may cross-react with Smad5 and Smad8 when phosphorylated at the equivalent sites. The antibody does not cross-react with Smad2.
Source / Purification	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser423 and Ser425 of human Smad3. Antibodies are purified by protein A and peptide affinity chromatography.
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). Following stimulation by TGF-β, Smad2 and Smad3 become phosphorylated at their carboxyl termini (Ser465 and 467 on Smad2; Ser423 and 425 on Smad3) by TGF-β Receptor I. Phosphorylated Smad 2/3 can complex with Smad4, translocate to the nucleus and regulate gene expression (9-11).
Background References	<ol style="list-style-type: none"> Heldin, C.H. et al. (1997) <i>Nature</i> 390, 465-71. Attisano, L. and Wrana, J.L. (1998) <i>Curr Opin Cell Biol</i> 10, 188-94. Derynck, R. et al. (1998) <i>Cell</i> 95, 737-40. Massagué, J. (1998) <i>Annu Rev Biochem</i> 67, 753-91. Whitman, M. (1998) <i>Genes Dev</i> 12, 2445-62. Wrana, J.L. (2000) <i>Sci STKE</i> 2000, re1. Attisano, L. and Wrana, J.L. (2002) <i>Science</i> 296, 1646-7. Moustakas, A. et al. (2001) <i>J Cell Sci</i> 114, 4359-69. Abdollah, S. et al. (1997) <i>J. Biol. Chem.</i> 272, 27678-27685. Souchelnytskyi, S. et al. (1997) <i>J. Biol. Chem.</i> 272, 28107-28115. Liu, X. et al. (1997) <i>Proc. Natl. Acad. Sci. USA</i> 94, 10669-10674.

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