[Primary Antibody]

phospho-Smad3 (Ser425) Rabbit pAb



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– DATASHEET –		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal	-	IHC-P (1:100-500)
GenelD: 4088	SWISS: P84022	IF (1:100-500)
Target: Smad3 (Ser425)		Flow-Cyt (1µg/Test)
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human Smad3 around the phosphorylation site of Ser425: SV(p-S)-NH2.		Reactivity: Human, Mouse, Rat (predicted: Rabbit, Pig, Cow, Chicken, Horse)
Purification: affinity purified by Protein A		
Concentration: 1mg/ml		Predicted MW.: 47 kDa
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		
		Subcellular Location: Cytoplasm ,Nucleus
Background: Smad3 is a 50 kDa me mediators of TGF bet differentiation and de three subclasses: rece receptor regulated (S (Smad 1, 5, and 8); the its interaction to the (Smad6 and 7). Activa TGF beta stimulation nucleus, allowing its b Phosphorylation of th 425 in the C terminus activity and TGF beta	mber of a family of proteins that act as key a superfamily signaling in cell proliferation, welopment. The Smad family is divided into eptor regulated Smads, activin/TGF beta mad2 and 3) or BMP receptor regulated e common partner, (Smad4) that functions via various Smads; and the inhibitory Smads, ted Smad3 oligomerizes with Smad4 upon and translocates as a complex into the binding to DNA and transcription factors. te two TGF beta dependent serines 423 and of Smad3 is critical for Smad3 transcriptional signaling.	

— VALIDATION IMAGES



Sample: Lane 1: Cerebrum (Mouse) Lysate at 40 ug Lane 2: Heart (Mouse) Lysate at 40 ug Lane 3: Kidney (Mouse) Lysate at 40 ug Lane 4: Cerebrum (Rat) Lysate at 40 ug Primary: Anti-Phospho-Smad3 (Ser425) (bs-5616R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 52 kD Observed band size: 50 kD



Paraformaldehyde-fixed, paraffin embedded (mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-Smad3 (Ser425)) Polyclonal Antibody, Unconjugated (bs-5616R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (rat brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-Smad3 (Ser425)) Polyclonal Antibody, Unconjugated (bs-5616R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control (Black line): HUVEC (Black).

Primary Antibody (green line): Rabbit Anti-Phospho-Smad3 (Ser425) antibody (bs-5616R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-AF647 Dilution: 1µg /test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

- [IF=4.545] Han X et al. The intervention effect of nicotine on cervical fibroblast-myofibroblast differentiation in lipopolysaccharide-induced preterm birth model through activating the TGF-β1/Smad3 pathwayBiomed Pharmacother.2020 Dec 24;134:111135. WB ;Mouse. 33352448
- [IF=4.271] Biao Li. et al. 20-Hydroxytetraenoic acid induces hepatic fibrosis via the TGF-β1/Smad3 signaling pathway. TOXICOL LETT. 2023 Jan;373:1 WB ;Human, Mouse. 36368619
- [IF=3.8] Tiancheng Lyu. et al.Naringin in repairing articular cartilage injury by activating TGF-β/Smad signaling pathway to attenuate inflammatory response..ARCHIVES OF BIOCHEMISTRY AND BIOPHYSICS.2025 Mar 20:768:110396. IF ;Rabbit. 40120921
- [IF=2.728] Xijuan Liu et al. Chondrocyte suppression is mediated by miR 129 5p via GDF11/SMAD3 signaling in developmental dysplasia of the hip. J Orthop Res. 2020 Dec;38(12):2559-2572. WB ;Rabbit. 32396235