

## **Product Data Sheet**

### **Anti-SMAD3 Antibody**

Catalog #	Source	e Reactivity	Applications	
CPA9444	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	o SMAD3	
Immunogen		Recombinant protein corresp	ponding to human SMAD3.	
Purification		The antibody was purified by	/ immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of SMAD3 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/1000 - 1/2000), IH (1/2	00 - 1/500)	
Gene Symbol		SMAD3		
Alternative Names		MADH3; Mothers against decapentaplegic homolog 3; MAD homolog 3; Mad3;		
		Mothers against DPP homolo	og 3; hMAD-3; JV15-2; SMAD family member 3; SMAD 3;	
		Smad3; hSMAD3		
Entrez Gene		4088 (Human)		
SwissProt		P84022 (Human)		
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
		freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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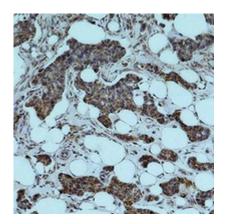
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**kDa** A B C D E F 200 140 100 80 60 50 40 30 20 For research purposes only, not for human use

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Western blot analysis of SMAD3 expression in A431 (A), Hela (B), C2C12 (C), NIH3T3 (D), PC12 (E), rat brain (F) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 50 kD)



Immunohistochemical analysis of SMAD3 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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