

# **Product Data Sheet**

### **Anti-SMAD3 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA5541	Rabbit	H, M, C	WB, IH		
Description	F	Rabbit polyclonal antibody t	o SMAD3		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
	r	region of human SMAD3. Th	e exact sequence is proprietary.		
Purification	٦	The antibody was purified b	/ immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of SMAD3 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	õ	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	9	SMAD3			
Alternative N	ames I	MADH3; Mothers against de	capentaplegic homolog 3; MAD homolog 3; Mad3;		
	ſ	Mothers against DPP homol	og 3; hMAD-3; JV15-2; SMAD family member 3; SMAD 3;		
	9	Smad3; hSMAD3			
Entrez Gene	2	4088 (Human); 17127 (Mou	se)		
SwissProt	F	P84022 (Human); Q8BUN5 (	Mouse)		
Storage/Stabi	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid				
	f	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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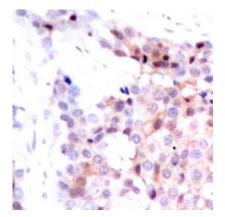
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**kDa** <u>A</u> <u>B</u> <u>C</u> <u>D</u> 150 100 70 50 35 25 Western blot analysis of SMAD3 expression in HEK293T (A), HeLa (B), A549 (C), mouse brain (D) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 52 kD)



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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