

Product Data Sheet

Anti-TEAD3 Antibody

Catalog #	Source	Reactivity	Applications
CQA3341	Rabbit	H, M, R	WB, IH, IF/IC
Description	Ral	bbit polyclonal antibody	to TEAD3
Immunogen	Red	combinant full length pro	tein of human TEAD3
Purification	The	e antibody was purified b	y immunogen affinity chromatography.
Specificity	Red	cognizes endogenous lev	els of TEAD3 protein.
Clonality	Pol	lyclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WE	3 (1/500 - 1/2000), IH (1/50) - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	TE	AD3	
Alternative Na	ames TEA	AD5; TEF5; Transcription	al enhancer factor TEF-5; DTEF-1; TEA domain family
	me	ember 3; TEAD-3	
Entrez Gene	700	05 (Human); 21678 (Mou	ise)
SwissProt	Q9	9594 (Human); P70210 (Mouse)
Storage/Stabi	lity Shi	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	fre	eze/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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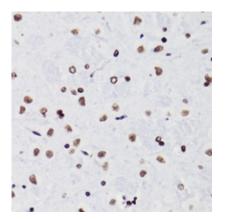
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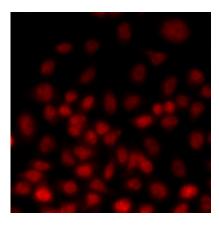
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Western blot analysis of TEAD3 expression in MCF7 (A), mouse heart (B), mouse brain (C) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 49 kD)



Immunohistochemical analysis of TEAD3 staining in human prostate 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.



Immunofluorescent analysis of TEAD3 staining in A549 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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