

## **Product Data Sheet**

### **Anti-TAO1 Antibody**

Catalog #	Source	e Reactivity	Applications		
CPA4843	Rabbit	t H, M, R	WB, IH		
Description		Rabbit polyclonal antibody	to TAO1		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human TAO1. The exact sequence is proprietary.			
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	els of TAO1 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 (1/500 - 1/1000), IH (1/1	00 - 1/200)		
Gene Symbol		TAOK1			
Alternative Names		KIAA1361; MAP3K16; MARKK; Serine/threonine-protein kinase TAO1; Kinase from			
		chicken homolog B; hKFC-B	MARK Kinase; MARKK; Prostate-derived sterile 20-like		
		kinase 2; PSK-2; PSK2; Prost	ate-derived STE20-like kinase 2; Thousand and one		
		amino acid protein kinase 1	; TAOK1; hTAOK1		
Entrez Gene		57551 (Human); 216965 (N	ouse); 286993 (Rat)		
SwissProt		Q7L7X3 (Human); Q5F2E8 (	Mouse); O88664 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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

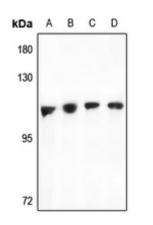
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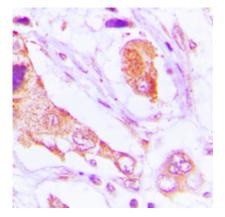


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Western blot analysis of TAO1 expression in HEK293T (A), A2780 (B), mouse testis (C), rat testis (D) whole cell lysates. (Predicted band size: 116 kD; Observed band size: 116 kD)



Immunohistochemical analysis of TAO1 staining in human lung 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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