

# **Product Data Sheet**

## Anti-S6K1 Antibody

Catalog #	Source	e Reactivity	Applications	
CPA9440	Rabbit	t H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	o S6K1	
Immunogen		Recombinant protein corres	oonding to human S6K1.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of S6K1 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium p	hosphate, 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		RPS6KB1		
Alternative Names		STK14A; Ribosomal protein S6 kinase beta-1; S6K-beta-1; S6K1; 70 kDa ribosomal		
		protein S6 kinase 1; P70S6K	L; p70-S6K 1; Ribosomal protein S6 kinase I;	
		Serine/threonine-protein kir	ase 14A; p70 ribosomal S6 kinase alpha; p70 S6 kinase	
		alpha; p70 S6K-alpha; p70 S	БКА	
Entrez Gene		6198 (Human)		
SwissProt		P23443 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	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

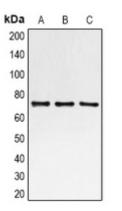
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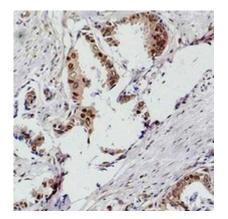


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Western blot analysis of S6K1 expression in Hela (A), NIH3T3 (B), PC12 (C) whole cell lysates. (Predicted band size: 59 kD; Observed band size: 70; 85 kD)



Immunohistochemical analysis of S6K1 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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