

## **Product Data Sheet**

### Anti-S6K1 (pT444) Antibody

Catalog #	Source	Reactivity	Applications		
CPA3314	Rabbit	H, M, R, B, C, D, P, Rb	WB, IH, IF/IC		
Description	R	Rabbit polyclonal antibody to S6K1 (pT444)			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
	re	region of human S6K1 (pT444). The exact sequence is proprietary.			
Purification	Т	he antibody was purified by imn	nunogen affinity chromatography.		
Specificity	R	ecognizes endogenous levels of	S6K1 (pT444) protein.		
Clonality	P	olyclonal			
Form	Li	iquid in 0.42% Potassium phospl	nate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	a	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/1000), IH (1/100 - 1/	200), IF/IC (1/100 - 1/500)		
Gene Symbol	R	PS6KB1			
Alternative Names		STK14A; Ribosomal protein S6 kinase beta-1; S6K-beta-1; S6K1; 70 kDa ribosomal			
	р	rotein S6 kinase 1; P70S6K1; p70	)-S6K 1; Ribosomal protein S6 kinase I;		
	S	erine/threonine-protein kinase	14A; p70 ribosomal S6 kinase alpha; p70 S6 kinase		
	a	lpha; p70 S6K-alpha;			
Entrez Gene		6198 (Human); 72508 (Mouse); 83840 (Rat)			
SwissProt	P	23443 (Human); Q8BSK8 (Mous	e); P67999 (Rat)		
Storage/Stabi	i <b>lity</b> Sl	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
	fr	reeze/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

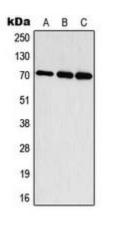
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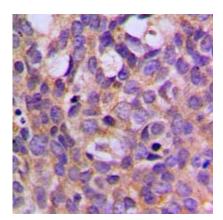
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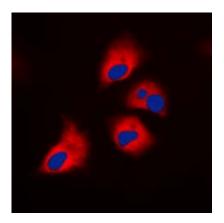
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Western blot analysis of S6K1 (pT444) expression in HepG2 EGF-treated (A), mouse brain (B), rat brain (C) whole cell lysates.



Immunohistochemical analysis of S6K1 (pT444) 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.



Immunofluorescent analysis of S6K1 (pT444) staining in HepG2 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. DAPI was used to stain the cell nuclei (blue).

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