

## **Product Data Sheet**

### Anti-SGK1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4230	Rabbit	H, M, R, B, C, Mk, Rb, Z	WB, IH, IF/IC		
Description	Rabbi	t polyclonal antibody to SGK1			
Immunogen	KLH-c	onjugated synthetic peptide e	ncompassing a sequence within the center		
	regio	n of human SGK1. The exact se	equence is proprietary.		
Purification	The a	The antibody was purified by immunogen affinity chromatography.			
Specificity	Recog	Recognizes endogenous levels of SGK1 protein.			
Clonality	Polyc	onal			
Form	Liquic	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 (1/100 - 1/20	0), IF/IC (1/100 - 1/500)		
Gene Symbol	SGK1				
Alternative Na	mes SGK; S	Serine/threonine-protein kina	se Sgk1; Serum/glucocorticoid-regulated kinase 1		
Entrez Gene	6446	(Human); 20393 (Mouse)			
SwissProt	0001	O00141 (Human); Q9WVC6 (Mouse); Q06226 (Rat)			
Storage/Stabili	<b>ty</b> Shipp	ed at 4°C. Upon delivery aliqu	ot and store at -20°C for one year. Avoid		
	freeze	e/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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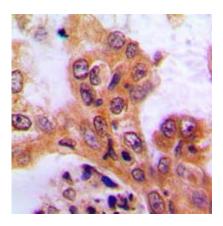
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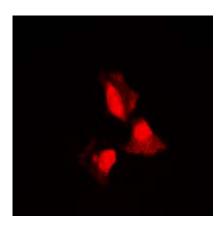
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Western blot analysis of SGK1 expression in Jurkat (A), HepG2 (B) whole cell lysates.



Immunohistochemical analysis of SGK1 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of SGK1 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 hidified 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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