

Product Data Sheet

Anti-GAK Antibody

Catalog #	Source	Reactivity	Applications		
CPA3952	Rabbit	н	WB, IH, IF/IC		
Description	Ra	abbit polyclonal antibody	r to GAK		
Immunogen	KL	H-conjugated synthetic	peptide encompassing a sequence within the N-term		
	re	gion of human GAK. The	exact sequence is proprietary.		
Purification	Th	ne antibody was purified	by immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous le	vels of GAK protein.		
Clonality	Pc	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	'B (1/200 - 1/500), IH (1/50) - 1/100), IF/IC (1/50 - 1/100)		
Gene Symbol	GA	AK			
Alternative Na	ames Cy	clin-G-associated kinase			
Entrez Gene 258		2580 (Human)			
SwissProt	01	14976 (Human)			
Storage/Stabi	lity Sh	nipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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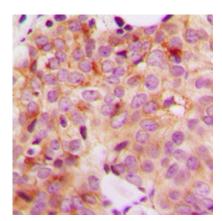
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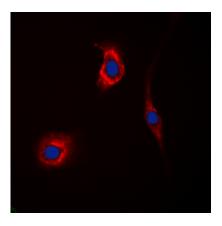
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Western blot analysis of GAK expression in PC3 (A), SGC7901 (B), DLD (C) whole cell lysates. (Predicted band size: 143 kD; Observed band size: 180 kD)



Immunohistochemical analysis of GAK 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 GAK staining in Hela 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. DAPI was used to stain the cell nuclei (blue).

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