

Product Data Sheet

Anti-SPINK6 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4254	Rabbit	H, M, R	WB, IH		
Description	Rat	Rabbit polyclonal antibody to SPINK6			
Immunogen	KLF	KLH-conjugated synthetic peptide encompassing a sequence within the center			
	reg	region of human SPINK6. The exact sequence is proprietary.			
Purification	The	e antibody was purified by	immunogen affinity chromatography.		
Specificity	Rec	cognizes endogenous leve	ls of SPINK6 protein.		
Clonality	Pol	yclonal			
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WB	8 (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	SPI	NK6			
Alternative Na	ames Ser	ine protease inhibitor Kaz	al-type 6; Kallikrein inhibitor		
Entrez Gene	404	1203 (Human); 433180 (N	louse); 100911369, 408235 (Rat)		
SwissProt	Q61	UWN8 (Human); Q8BT20	(Mouse); Q6IE47 (Rat)		
Storage/Stabi	lity Shi	pped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	free	eze/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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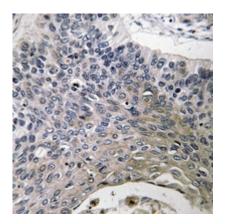
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A B C

For research purposes only, not for human use

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Western blot analysis of SPINK6 expression in HeLa (A), Raw264.7 (B), H9C2 (C) whole cell lysates. (Predicted band size: 8 kD; Observed band size: 8 kD)



Immunohistochemical analysis of SPINK6 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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