

Product Data Sheet

Anti-VRK3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5818	Rabbit	H <i>,</i> R	WB, IH		
Description		Rabbit polyclonal antibody	to VRK3		
Immunogen		KLH-conjugated synthetic po	eptide encompassing a sequence within the center		
		region of human VRK3. The	exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of VRK3 protein.		
Clonality		Polyclonal			
Conjugation					
Form		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 (1/50) - 1/100)		
Gene Symbol		VRK3			
Alternative Na	ames	Inactive serine/threonine-p	rotein kinase VRK3; Serine/threonine-protein		
		pseudokinase VRK3; Vaccini	a-related kinase 3		
Entrez Gene		51231 (Human)			
SwissProt		Q8IV63 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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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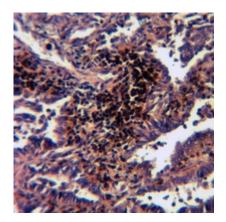


For research purposes only, not for human use

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KDa A B C D 70 50 35 25 20

Western blot analysis of VRK3 expression in HEK293T (A), A549 (B), U87MG (C), rat liver (D) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 53 kD)



Immunohistochemical analysis of VRK3 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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