

Product Data Sheet

Anti-AKT2 (pS474) Antibody

Catalog #	Source	Reactivity	Applications			
CPA7122	Rabbit	H, M, R, C, D	WB, IH			
Description		Rabbit polyclonal antibody to AKT2 (pS474)				
Immunogen		KLH-conjugated synthetic pepti	de encompassing a sequence within the C-term			
		region of human AKT2 (pS474).	The exact sequence is proprietary.			
Purification		The antibody was purified by ir	nmunogen affinity chromatography.			
Specificity		Recognizes endogenous levels	of AKT2 (pS474) protein.			
Clonality		Polyclonal				
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 (1/100 -	1/200)			
Gene Symbol		AKT2				
Alternative N	ames	RAC-beta serine/threonine-pro	tein kinase; Protein kinase Akt-2; Protein kinase B			
		beta; PKB beta; RAC-PK-beta				
Entrez Gene		208 (Human); 11652 (Mouse); 25233 (Rat)				
SwissProt		P31751 (Human); Q60823 (Mouse); P47197 (Rat)				
Storage/Stabi	ility	Shipped at 4°C. Upon delivery a	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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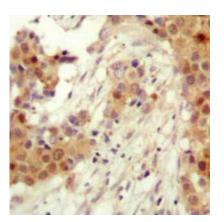
200

kDa A B C

For research purposes only, not for human use

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Western blot analysis of AKT2 (pS474) expression in Hela (A), mouse brain (B), PC12 UV-treated (C) whole cell lysates.



Immunohistochemical analysis of AKT2 (pS474) 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 conjugacompact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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