

Product Data Sheet

Anti-Annexin A3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1051	Rabbit	H, M, R, D, Mk, Rb	WB, IH		
Description	Ra	abbit polyclonal antibody to A	Annexin A3		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
	re	region of human Annexin A3. The exact sequence is proprietary.			
Purification	Tł	he antibody was purified by ir	nmunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous levels	of Annexin A3 protein.		
Clonality	Pc	olyclonal			
Conjugation					
Form	Lie	quid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/100 -	1/200)		
Gene Symbol	A	NXA3			
Alternative Na	ames Al	NX3; Annexin A3; 35-alpha ca	lcimedin; Annexin III; Annexin-3; Inositol 1,2-cyclic		
	pł	hosphate 2-phosphohydrolas	e; Lipocortin III; Placental anticoagulant protein III;		
	PA	AP-III			
Entrez Gene	30	06 (Human); 11745 (Mouse);	25291 (Rat)		
SwissProt	P1	12429 (Human); O35639 (Mo	use); P14669 (Rat)		
Storage/Stabi	lity Sh	nipped at 4°C. Upon delivery a	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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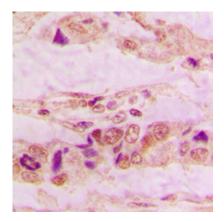
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Western blot analysis of Annexin A3 expression in Hela (A) whole cell lysates. (Predicted band size: 36 kD; Observed band size: 36 kD)



Immunohistochemical analysis of Annexin A3 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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