

Product Data Sheet

Anti-VASP (Phospho-S157) Antibody

Catalog #	Source	Reactivity	Applications		
CPA3352	Rabbit	H <i>,</i> R	WB, IH		
Description	Ra	Rabbit polyclonal antibody to VASP (Phospho-S157)			
Immunogen	KL	LH-conjugated synthetic pl	nosphopeptide corresponding to residues surrounding		
	S1	157 of human VASP protei	n. The exact sequence is proprietary.		
Purification	Tł	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of VASP protein only when phosphorylated at S157.		
Clonality	Pc	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	VA	ASP			
Alternative Na	ames Va	asodilator-stimulated phos	phoprotein; VASP		
Entrez Gene	74	408 (Human)			
SwissProt	PS	50552 (Human)			
Storage/Stabi	lity Sł	nipped at 4°C. Upon delive	ry 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

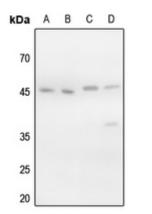
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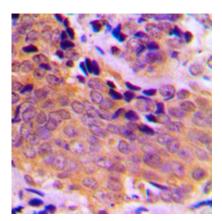


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Western blot analysis of VASP (Phospho-S157) expression in HEK293T (A), Hela (B), rat lung (C), rat spleen (D) whole cell lysates. (Predicted band size: 39 kD; Observed band size: 50 kD)



Immunohistochemical analysis of VASP (Phospho-S157) 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.

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