

Product Data Sheet

Anti-p47 phox (Phospho-S345) Antibody

Catalog #	Source	Reactivity	Applications		
CPA5058	Rabbit	H <i>,</i> M <i>,</i> R	WB, IH		
Description		Rabbit polyclonal antibody to	o p47 phox (Phospho-S345)		
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
		S345 of human p47 phox pro	tein. The exact sequence is proprietary.		
Purification		The antibody was purified by	r immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of p47 phox protein only when phosphorylated at			
		S345.			
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol		NCF1			
Alternative Names		NOXO2; SH3PXD1A; Neutrophil cytosol factor 1; NCF-1; 47 kDa autosomal chronic			
		granulomatous disease prote	ein; 47 kDa neutrophil oxidase factor; NCF-47K;		
		Neutrophil NADPH oxidase f	actor 1; Nox organizer 2; Nox-organizing protein 2; SH3		
		and PX domain-contain			
Entrez Gene		653361 (Human); 17969 (Mo	ouse)		
SwissProt		P14598 (Human); Q09014 (N	1ouse)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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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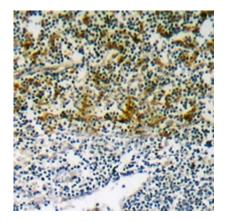
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Western blot analysis of p47 phox (Phospho-S345) expression in rat lung (A), rat liver (B) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 47 kD)



Immunohistochemical analysis of p47 phox (Phospho-S345) staining in human lymph node 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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