

**p47-phox (phospho Ser328) Polyclonal Antibody**

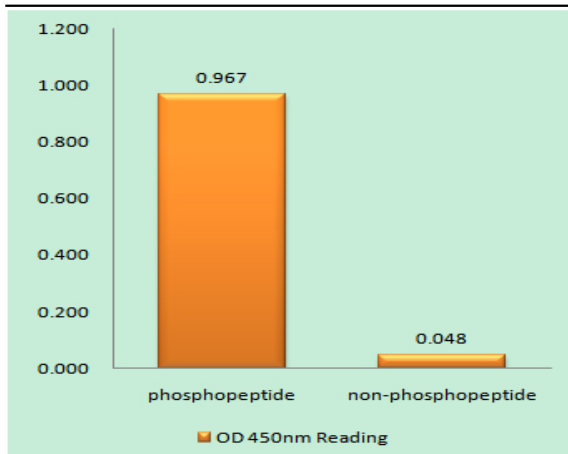
<b>Catalog No :</b>	YP1018
<b>Reactivity :</b>	Human;Mouse;Rat;Cow
<b>Applications :</b>	WB;IHC;IF;ELISA
<b>Target :</b>	p47-phox
<b>Fields :</b>	>>Chemokine signaling pathway;>>Phagosome;>>Osteoclast differentiation;>>Neutrophil extracellular trap formation;>>Fc gamma R-mediated phagocytosis;>>Leukocyte transendothelial migration;>>Prion disease;>>Leishmaniasis;>>Chemical carcinogenesis - reactive oxygen species;>>Diabetic cardiomyopathy;>>Lipid and atherosclerosis;>>Fluid shear stress and atherosclerosis
<b>Gene Name :</b>	NCF1
<b>Protein Name :</b>	Neutrophil cytosol factor 1
<b>Human Gene Id :</b>	653361
<b>Human Swiss Prot No :</b>	P14598
<b>Mouse Gene Id :</b>	17969
<b>Mouse Swiss Prot No :</b>	Q09014
<b>Immunogen :</b>	The antiserum was produced against synthesized peptide derived from human Neutrophil Cytosol Factor 1 around the phosphorylation site of Ser328. AA range:301-350
<b>Specificity :</b>	Phospho-p47-phox (S328) Polyclonal Antibody detects endogenous levels of p47-phox protein only when phosphorylated at S328.
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Polyclonal, Rabbit,IgG
	WB 1:500-2000 IHC 1:100 - 1:300. ELISA: 1:10000. IF 1:50-200

---

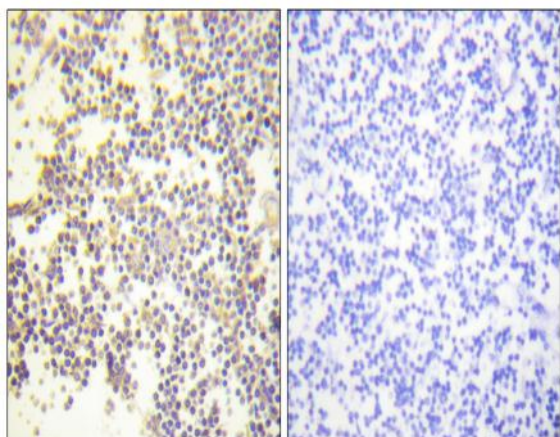
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Concentration :</b>	1 mg/ml
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)
<b>Molecularweight :</b>	45kD
<b>Cell Pathway :</b>	Chemokine;Fc gamma R-mediated phagocytosis;Leukocyte transendothelial migration;
<b>Background :</b>	The protein encoded by this gene is a 47 kDa cytosolic subunit of neutrophil NADPH oxidase. This oxidase is a multicomponent enzyme that is activated to produce superoxide anion. Mutations in this gene have been associated with chronic granulomatous disease. [provided by RefSeq, Jul 2008],
<b>Function :</b>	disease:Defects in NCF1 are the cause of chronic granulomatous disease autosomal recessive cytochrome-b-positive type 1 (CGD1) [MIM:233700]. Chronic granulomatous disease is a genetically heterogeneous disorder characterized by the inability of neutrophils and phagocytes to kill microbes that they have ingested. Patients suffer from life-threatening bacterial/fungal infections.,function:NCF2, NCF1, and a membrane bound cytochrome b558 are required for activation of the latent NADPH oxidase (necessary for superoxide production).,online information:NCF1 deficiency database,similarity:Contains 1 PX (phox homology) domain.,similarity:Contains 2 SH3 domains.,subunit:Interacts with NOXA1.,
<b>Subcellular Location :</b>	Cytoplasm, cytosol . Membrane ; Peripheral membrane protein ; Cytoplasmic side .
<b>Expression :</b>	Detected in peripheral blood monocytes and neutrophils (at protein level).

---

## Products Images



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using Neutrophil Cytosol Factor 1 (Phospho-Ser328) Antibody



Immunohistochemistry analysis of paraffin-embedded human tonsil, using Neutrophil Cytosol Factor 1 (Phospho-Ser328) Antibody. The picture on the right is blocked with the phosphopeptide.