#### Anti-p47 phox Rabbit Polyclonal Antibody



# **Catalog #: 51671**

#### **Aliases**

NOXO2; SH3PXD1A; Neutrophil cytosol factor 1; NCF-1; 47 kDa autosomal chronic granulomatous disease protein; 47 kDa neutrophil oxidase factor; NCF-47K; Neutrophil NADPH oxidase factor 1; Nox organizer 2; Nox-organizing protein 2; SH3 and PX domain-containing protein 1A; p47-phox

# **Background**

Gene Name: NCF1

NCBI Gene Entry: 653361 UniProt Entry: P14598

# **Application Information**

Molecular Weight: Predicted, 44 kDa; observed, 47 kDa

Clonality: Rabbit polyclonal antibody

Species Reactivity: Human, mouse, rat, bovine, dog, monkey, sheep

Applications Tested: Western blotting (WB), immunohistochemistry (IHC), immunocytochemistry

(IC)

#### **Immunogen**

A synthesized peptide derived from human p47 phox

# **Isotype**

Rabbit IgG

#### **Storage Buffer**

Supplied in PBS (pH 7.3) containing 30% glycerol, and 0.01% sodium azide.

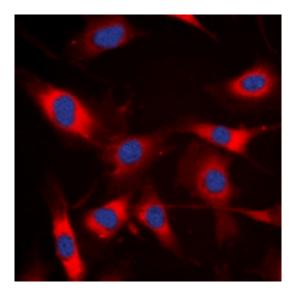
#### **Storage**

Store at -20 °C for one year.

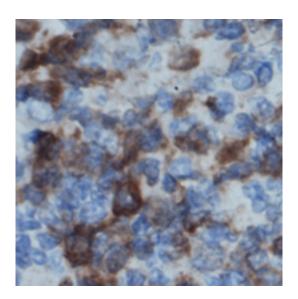
#### **Recommended Dilutions**

Western Blotting (WB): 1:500-1:1,000 Immunohistochemistry (IHC): 1:100-1:200 Immunocytochemistry (IC): 1:100-1:500 **Note:** This product is for research use only.

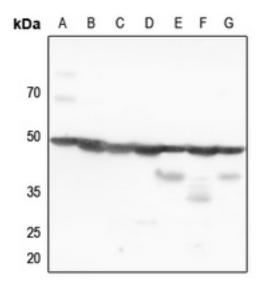
#### **Validation Data**



Immunocytochemical analysis of p47 phox staining in Raji cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).



Immunohistochemical analysis of p47 phox staining in human lymphoma 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.



Western blotting analysis of p47 phox expression in HEK293T (A), Hela (B), mouse lung (C), mouse liver (D), rat lung (E), rat liver (F), rat spleen (G) whole cell lysates.