

PRODUCT DATA SHEET

Bioworld Technology,Inc.

p47-phox (phospho-S370) polyclonal antibody

Catalog: BS4852 Host: Rabbit Reactivity: Human, Mouse, Rat

BackGround:

The heredity disease chronic granulomatous disease (CGF) has been linked to mutations in p47-phox and p67-phox. The cytosolic proteins p47-phox p67-phox, also designated neutrophil cytosol factor (NCF)1 and NCF2, respectively, are required for activation of the superoxide-producing NADPH oxidase in neutrophils and other phagocytic cells. During activation of the NADPH oxidase, p47-phox and p67-phox migrate to the plasma membrane where they associate with cytochrome b558 and the small G protein Rac to form the functional enzyme complex. Both p47-phox and p67-phox contain two Src homology 3 (SH3) domains. The C-terminal SH3 doamin of p67-phox has been shown to interact with the proline rich domain of p47-phox, suggesting that p47-phox may faciliate the transport of p67-phox to the membrane.

Product:

1 mg/ml in Phosphate buffered saline (PBS) with 0.05% sodium azide, approx. pH 7.2.

Molecular Weight:

~ 45 kDa

Swiss-Prot:

P14598

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB: 1:500~1:1000 IHC: 1:50~1:200 ICC: 1:50~1:200 IP: 1:50~1:200

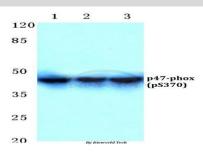
Storage&Stability:

Store at $4 \,\mathrm{C}$ short term. Aliquot and store at $-20 \,\mathrm{C}$ long term. Avoid freeze-thaw cycles.

Specificity:

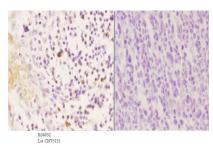
p-p47-phox (S370) polyclonal antibody detects endogenous levels of p47-phox protein only when phosphorylated at Ser370.

DATA:



Western blot (WB) analysis of p-p47-phox (S370) polyclonal antibody at 1:500 dilution

Lane1:Hela cell treated with serum starvation(24h)
Lane2:Raw264.7 cell treated with serum starvation(24h)
Lane3:H9C2 cell treated with serum starvation(24h)



Immunohistochemistry (IHC) analyzes of p-p47-phox (S370) pAb in paraffin-embedded human tonsil carcinoma tissue at 1:50.showing cytoplasmic and nucleus staining. Negative control (the right)Using PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG-biotin followed by avidin-peroxidase.

Note:

For research use only, not for use in diagnostic procedure.

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