

Technical Data Sheet

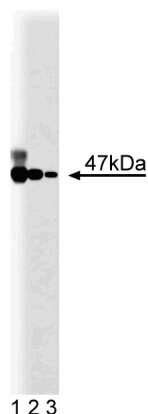
Purified Mouse Anti-Human p47[phox]**Product Information**

Material Number:	610355
Size:	150 µg
Concentration:	250 µg/ml
Clone:	1/p47Phox
Immunogen:	Human p47[phox] aa. 18-197
Isotype:	Mouse IgG1
Reactivity:	QC Testing: Human
Target MW:	47 kDa
Storage Buffer:	Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium azide.

Description

The neutrophil respiratory burst oxidase (NADPH-oxidase) generates superoxide and secondary oxygen-derived toxic products in response to bacteria or a variety of soluble stimuli. The enzyme is dormant in resting neutrophils. The active site of this enzyme is located in an integral membrane cytochrome, b558, which consists of the two subunits gp91[phox] and p21[phox]. Superoxide production depends on the formation of a complex that includes two cytosolic proteins, p67[phox] and p47[phox]. The GTP-binding protein Rac is also an essential component for oxidase activity. p47[phox] is a highly basic protein that contains two SH3 domains. The C-terminal quarter of the molecule contains many potential phosphorylation sites, consisting of serines and basic residues. Expression of p47[phox] is restricted to cells of phagocytic or lymphocytic lineage. IFN-γ is a potent inducer of both p47[phox] mRNA and protein. p47[phox] is an early reactant in oxidase assembly and this assembly can be inhibited by a C-terminal peptide of the large subunit of cytochrome b558. It is thought that p47[phox] binds directly to the cytochrome, while p67[phox] associates with the cytochrome by binding p47[phox].

This antibody is routinely tested by western blot analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.



Western blot analysis of p47[phox] on EB-1 lysate. 1:500 (lane 1), 1:1000 (lane2), 1:2000 (lane 3) dilution of anti-p47[phox] antibody.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

Store undiluted at -20° C.

Application Notes**Application**

Western blot	Routinely Tested
Immunofluorescence	Tested During Development

BD Biosciences

bdbiosciences.com

United States	Canada	Europe	Japan	Asia Pacific	Latin America/Caribbean
877.232.8995	888.259.0187	32.53.720.550	0120.8555.90	65.6861.0633	55.11.5185.9995

For country-specific contact information, visit bdbiosciences.com/how_to_order/

Conditions: The information disclosed herein is not to be construed as a recommendation to use the above product in violation of any patents. BD Biosciences will not be held responsible for patent infringement or other violations that may occur with the use of our products. Purchase does not include or carry any right to resell or transfer this product either as a stand-alone product or as a component of another product. Any use of this product other than the permitted use without the express written authorization of Becton Dickinson and Company is strictly prohibited.

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Not for resale.

BD, BD Logo and all other trademarks are the property of Becton, Dickinson and Company. ©2006 BD



Suggested Companion Products

<u>Catalog Number</u>	<u>Name</u>	<u>Size</u>	<u>Clone</u>
554002	HRP Goat Anti-Mouse Igs	1.0 ml	(none)

Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to www.bdbiosciences.com/pharming/protocols for technical protocols.
3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

References

- Ago T, Nunoi H, Ito T, Sumimoto H. Mechanism for phosphorylation-induced activation of the phagocyte NADPH oxidase protein p47(phox). Triple replacement of serines 303, 304, and 328 with aspartates disrupts the SH3 domain-mediated intramolecular interaction in p47(phox), thereby activating the oxidase. *J Biol Chem.* 1999; 274(47):33644-33653.(Clone-specific: Western blot)
- Chanock SJ, el Benna J, Smith RM, Babior BM. The respiratory burst oxidase. *J Biol Chem.* 1994; 269(40):24519-24522.(Biology)
- Hata K, Ito T, Takeshige K, Sumimoto H. Anionic amphiphile-independent activation of the phagocyte NADPH oxidase in a cell-free system by p47phox and p67phox, both in C terminally truncated forms. Implication for regulatory Src homology 3 domain-mediated interactions. *J Biol Chem.* 1998; 273(7):4232-4236. (Biology)
- Jackson SH, Malech HL, Kozak CA, Lomax KJ, Gallin JI, Holland SM. Cloning and functional expression of the mouse homologue of p47phox. *Immunogenetics.* 1994; 39(4):272-275.(Biology)
- Shiose A, Sumimoto H. Arachidonic acid and phosphorylation synergistically induce a conformational change of p47phox to activate the phagocyte NADPH oxidase. *J Biol Chem.* 2000; 275(18):13793-13801.(Clone-specific: Western blot)