

# **NQO1** Antibody

Purified Mouse Monoclonal Antibody Catalog # AO1643a

## **Product Information**

**Application** WB, FC, E **Primary Accession** P15559 Reactivity Human Host Mouse Clonality Monoclonal **Clone Names** 1A11 Isotype IgG1 30868 **Calculated MW** 

**Description** This gene is a member of the NAD(P)H dehydrogenase (quinone) family and

encodes a cytoplasmic 2-electron reductase. This FAD-binding protein forms homodimers and reduces quinones to hydroquinones. This protein's enzymatic activity prevents the one electron reduction of quinones that results in the production of radical species. Mutations in this gene have been associated with tardive dyskinesia (TD), an increased risk of hematotoxicity after exposure to benzene, and susceptibility to various forms of cancer. Altered expression of this protein has been seen in many tumors and is also associated with Alzheimer's disease (AD). Alternate transcriptional splice

variants, encoding different isoforms, have been characterized.

**Immunogen** Purified recombinant fragment of human NQO1 expressed in E. Coli.

**Formulation** Ascitic fluid containing 0.03% sodium azide.

### **Additional Information**

**Gene ID** 1728

Other Names NAD(P)H dehydrogenase [quinone] 1, 1.6.5.2, Azoreductase, DT-diaphorase,

DTD, Menadione reductase, NAD(P)H:quinone oxidoreductase 1,

Phylloquinone reductase, Quinone reductase 1, QR1, NQO1, DIA4, NMOR1

**Dilution** WB~~1/500 - 1/2000 FC~~1/200 - 1/400 E~~1/10000

**Storage** Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** NQO1 Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

## **Protein Information**

Name

NQO1 {ECO:0000303 | PubMed:1657151, ECO:0000312 | HGNC:HGNC:2874}

**Function** 

Flavin-containing quinone reductase that catalyzes two- electron reduction of quinones to hydroquinones using either NADH or NADPH as electron donors. In a ping-pong kinetic mechanism, the electrons are sequentially transferred from NAD(P)H to flavin cofactor and then from reduced flavin to the guinone, bypassing the formation of semiguinone and reactive oxygen species (By similarity) (PubMed:8999809, PubMed:9271353). Regulates cellular redox state primarily through quinone detoxification. Reduces components of plasma membrane redox system such as coenzyme Q and vitamin quinones, producing antioxidant hydroquinone forms. In the process may function as superoxide scavenger to prevent hydroquinone oxidation and facilitate excretion (PubMed:15102952, PubMed:8999809, PubMed: 9271353). Alternatively, can activate quinones and their derivatives by generating redox reactive hydroquinones with DNA cross-linking antitumor potential (PubMed:8999809). Acts as a gatekeeper of the core 20S proteasome known to degrade proteins with unstructured regions. Upon oxidative stress, interacts with tumor suppressors TP53 and TP73 in a NADH-dependent way and inhibits their ubiquitin-independent degradation by the 20S proteasome (PubMed: 15687255, PubMed: 28291250).

**Cellular Location** 

Cytoplasm, cytosol {ECO:0000250|UniProtKB:P05982}

### References

1. Mol Cancer Ther. 2009 Dec;8(12):3369-78. 2. J Biol Chem. 2009 Nov 27;284(48):33233-41.

## **Images**

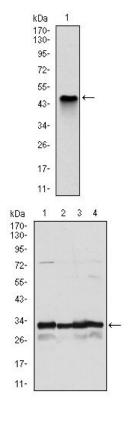
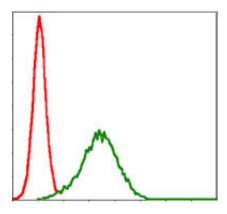


Figure 1: Western blot analysis using NQO1 mAb against human NQO1 (AA: 134-274) recombinant protein. (Expected MW is 41.3 kDa)

Figure 2: Western blot analysis using NQO1 mouse mAb against A549 (1), Hela (2), MCF-7 (3) and HepG2 (4) cell lysate.

Figure 4: Flow cytometric analysis of HepG2 cells using NQO1 mouse mAb (green) and negative control (red).



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