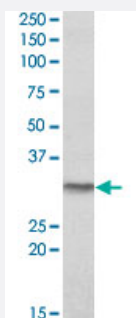


NQO1 (isoform A) polyclonal antibody

Catalog # PAB19680

Size 100 ug

Applications



Western Blot (Tissue lysate)

NQO1 (isoform A) polyclonal antibody (Cat # PAB19680) (0.3 ug/mL) staining of human kidney lysate (35 ug protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

Specification

Product Description	Goat polyclonal antibody raised against synthetic peptide of NQO1.
Immunogen	A synthetic peptide corresponding to internal region of human NQO1.
Sequence	C-DKGPFRRSKKAVLS
Host	Goat
Theoretical MW (kDa)	30.9
Reactivity	Human
Specificity	This antibody is expected to recognize reported isoform a (NP_000894.1) only.
Form	Liquid
Purification	Antigen affinity purification
Recommend Usage	ELISA (1:32000) Western Blot (0.3-1 ug/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	In Tris saline, pH 7.3 (0.5% BSA, 0.02% sodium azide)

Storage Instruction

Store at -20°C.
Aliquot to avoid repeated freezing and thawing.

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Western Blot (Tissue lysate)

NQO1 (isoform A) polyclonal antibody (Cat # PAB19680) (0.3 ug/mL) staining of human kidney lysate (35 ug protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

- Enzyme-linked Immunoabsorbent Assay

Gene Info — NQO1

Entrez GeneID[1728](#)**Protein Accession#**[NP_000894.1](#)**Gene Name**

NQO1

Gene Alias

DHQU, DIA4, DTD, NMOR1, NMORI, QR1

Gene Description

NAD(P)H dehydrogenase, quinone 1

Omim ID[125860](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

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. [provided by RefSeq]

Other Designations

DT-diaphorase|NAD(P)H menadione oxidoreductase 1, dioxin-inducible|NAD(P)H:Quinone acceptor oxidoreductase type 1|NAD(P)H:menadione oxidoreductase 1|NAD(P)H:quinone oxidoreductase|azoreductase|diaphorase (NADH/NADPH) (cytochrome b-5 reductase)|diaphorase-4|

Disease

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- [Acute Disease](#)
- [Acute Lung Injury](#)
- [Adenocarcinoma](#)
- [Adenoma](#)
- [Akathisia](#)
- [Alcohol Withdrawal Delirium](#)
- [Alzheimer disease](#)
- [Anemia](#)
- [Aneuploidy](#)
- [Angina Pectoris](#)
- [Asthma](#)
- [Atherosclerosis](#)
- [Atrophy](#)
- [Balkan Nephropathy](#)
- [Barrett Esophagus](#)
- [Bipolar Disorder](#)
- [Brain Neoplasms](#)
- [Breast cancer](#)
- [Breast Neoplasms](#)
- [Bronchial Hyperreactivity](#)
- [Burkitt lymphoma](#)
- [Carcinoma](#)

- [Cardiovascular Diseases](#)
- [Carotid Stenosis](#)
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- [Central Nervous System Diseases](#)
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- [Chromosome Deletion](#)
- [Chronic Disease](#)
- [Cleft Lip](#)
- [Cleft Palate](#)
- [Cocarcinogenesis](#)
- [Cognition Disorders](#)
- [Colitis](#)
- [Colon cancer](#)
- [Colonic Neoplasms](#)
- [Colorectal Neoplasms](#)
- [Coronary Artery Disease](#)
- [Coronary Disease](#)
- [Cystic fibrosis](#)
- [Depressive Disorder](#)
- [Diabetes Mellitus](#)
- [Diabetic Nephropathies](#)
- [Disease Progression](#)
- [Disease Susceptibility](#)
- [DNA Damage](#)
- [Drug Hypersensitivity](#)
- [Drug Toxicity](#)

- [Ductus Arteriosus](#)
- [Dyskinesia](#)
- [Dyskinesias](#)
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- [Hepatitis](#)
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- [Infertility](#)
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- [Kidney Failure](#)
- [Laryngeal Neoplasms](#)
- [Leber hereditary optic neuropathy](#)
- [Leukemia](#)
- [Liver Neoplasms](#)

- [Lung carcinoma](#)
- [Lung Neoplasms](#)
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- [Varicocele](#)
- [Werner syndrome](#)