

Anti-NDUFA8 Antibody

Rabbit polyclonal antibody to NDUFA8 Catalog # AP59632

Product Information

Application WB, IP, IHC
Primary Accession P51970
Other Accession O9DCI5

Reactivity Human, Mouse, Rat, Monkey, Bovine

Host Rabbit
Clonality Polyclonal
Calculated MW 20105

Additional Information

Gene ID 4702

Other Names NADH dehydrogenase [ubiquinone] 1 alpha subcomplex subunit 8; Complex

I-19kD; CI-19kD; Complex I-PGIV; CI-PGIV; NADH-ubiquinone oxidoreductase

19 kDa subunit

Target/Specificity Recognizes endogenous levels of NDUFA8 protein.

Dilution WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IP (1/10 - 1/100) IP~~N/A

IHC~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IP (1/10 - 1/100)

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name NDUFA8

Function Accessory subunit of the mitochondrial membrane respiratory chain NADH

dehydrogenase (Complex I), that is believed not to be involved in catalysis (PubMed:<u>27626371</u>, PubMed:<u>32385911</u>, PubMed:<u>33153867</u>). Complex I functions in the transfer of electrons from NADH to the respiratory chain (PubMed:<u>27626371</u>). The immediate electron acceptor for the enzyme is

believed to be ubiquinone (PubMed:27626371).

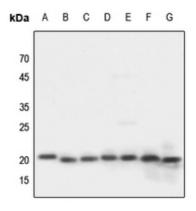
Cellular Location Mitochondrion inner membrane; Peripheral membrane protein.

Mitochondrion intermembrane space. Mitochondrion

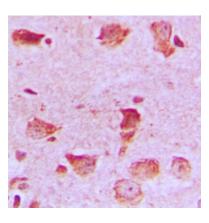
Background

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human NDUFA8. The exact sequence is proprietary.

Images



Western blot analysis of NDUFA8 expression in HEK293T (A), Hela (B), A2780 (C), HepG2 (D), mouse muscle (E), mouse kidney (F), rat kidney (G) whole cell lysates.



Immunohistochemical analysis of NDUFA8 staining in human brain 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.

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