NDUFB6 (B-2): sc-515596



The Power to Question

BACKGROUND

NDUFB6 (NADH dehydrogenase [ubiquinone] 1 β subcomplex subunit 6), also known as CI-B17 (complex I-B17) or NADH-ubiquinone oxidoreductase B17 subunit, is a 128 amino acid single-pass membrane protein that localizes to the matrix side of the mitochondrial inner membrane. A member of the complex I NDUFB6 subunit family, NDUFB6 is encoded by a gene that maps to human chromosome 9p21.1 and mouse chromosome 4 A5. Human chromosome 9 houses over 900 genes and comprises nearly 4% of the human genome. Hereditary hemorrhagic telangiectasia, which is characterized by harmful vascular defects, and familial dysautonomia, are both associated with chromosome 9. Notably, chromosome 9 encompasses the largest interferon family gene cluster.

REFERENCES

- Smeitink, J., et al. 1998. Molecular characterization and mutational analysis
 of the human B17 subunit of the mitochondrial respiratory chain complex I.
 Hum. Genet. 103: 245-250.
- 2. Murray, J., et al. 2003. The subunit composition of the human NADH dehydrogenase obtained by rapid one-step immunopurification. J. Biol. Chem. 278: 13619-13622.
- Zhuang, H., et al. 2006. Lupus-like disease and high interferon levels corresponding to trisomy of the type I interferon cluster on chromosome 9p. Arthritis Rheum. 54: 1573-1579.
- Burmeister, T., et al. 2007. Atypical Bcr-Abl mRNA transcripts in adult acute lymphoblastic leukemia. Haematologica 92: 1699-1702.

CHROMOSOMAL LOCATION

Genetic locus: NDUFB6 (human) mapping to 9p21.1.

SOURCE

NDUFB6 (B-2) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 99-115 near the C-terminus of NDUFB6 of human origin.

PRODUCT

Each vial contains 200 μ g IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

NDUFB6 (B-2) is available conjugated to agarose (sc-515596 AC), 500 $\mu g/0.25$ ml agarose in 1 ml, for IP; to HRP (sc-515596 HRP), 200 $\mu g/ml$, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-515596 PE), fluorescein (sc-515596 FITC), Alexa Fluor* 488 (sc-515596 AF488), Alexa Fluor* 546 (sc-515596 AF546), Alexa Fluor* 594 (sc-515596 AF594) or Alexa Fluor* 647 (sc-515596 AF647), 200 $\mu g/ml$, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor* 680 (sc-515596 AF680) or Alexa Fluor* 790 (sc-515596 AF790), 200 $\mu g/ml$, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-515596 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

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APPLICATIONS

NDUFB6 (B-2) is recommended for detection of NDUFB6 of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000)

Suitable for use as control antibody for NDUFB6 siRNA (h): sc-92468, NDUFB6 shRNA Plasmid (h): sc-92468-SH and NDUFB6 shRNA (h) Lentiviral Particles: sc-92468-V.

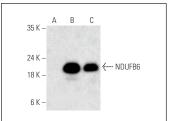
Molecular Weight of NDUFB6: 15 kDa.

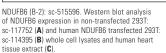
Positive Controls: human heart extract: sc-363763, Caki-1 cell lysate: sc-2224 or NDUFB6 (h): 293T Lysate: sc-114395.

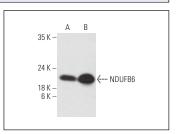
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz* Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz* Mounting Medium: sc-24941 or UltraCruz* Hard-set Mounting Medium: sc-359850.

DATA







NDUFB6 (B-2): sc-515596. Western blot analysis of NDUFB6 expression in Caki-1 whole cell lysate ($\bf A$) and human kidney tissue extract ($\bf B$).

SELECT PRODUCT CITATIONS

 Albanesi, J., et al. 2020. Transcriptional and metabolic dissection of ATRA-induced granulocytic differentiation in NB4 acute promyelocytic leukemia cells. Cells 9: 2423.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

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