abcam

Product datasheet

Anti-NDUFV1 antibody ab55535

3 References　画像数 2

製品の概要

製品名	Anti-NDUFV1 antibody
製品の詳細	Mouse monoclonal to NDUFV1
由来種	Mouse
アプリケーション	**適用あり:** WB, Flow Cyt
種交差性	**交差種:** Human
免疫原	Recombinant fragment: KAIARLIEFY KHESCGQCTP CREGVDWMNK VMARFVRGDA RPAEIDSLWE ISKQIEGHTI CALGDGAAWP VQGLIRHFRP ELEERMQRFA QQHQARQAAS , corresponding to amino acids 365-465 of Human NDUFV1 Run BLAST with ExPASy　Run BLAST with NCBI

製品の特性

製品の状態	Liquid
保存方法	Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.
バッファー	Preservative: None PBS, pH 7.2
精製度	Protein G purified
ポリ/モノ	**モノクローナル**
アイソタイプ	IgG2b
軽鎖の種類	kappa

アプリケーション

Our Abpromise guarantee covers the use of **ab55535** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

アプリケーション	Abreviews	特記事項
WB		Use a concentration of 1 - 5 µg/ml. Predicted molecular weight: 51 kDa.

アプリケーション	Abreviews	特記事項
Flow Cyt		Use 0.1µg for 10^6 cells. ab170192 - Mouse monoclonal IgG2b, is suitable for use as an isotype control with this antibody.

ターゲット情報

機能

Core subunit of the mitochondrial membrane respiratory chain NADH dehydrogenase (Complex I) that is believed to belong to the minimal assembly required for catalysis. Complex I functions in the transfer of electrons from NADH to the respiratory chain. The immediate electron acceptor for the enzyme is believed to be ubiquinone.

関連疾患

Defects in NDUFV1 are a cause of Leigh syndrome (LS) [MIM:256000]. LS is a severe neurological disorder characterized by bilaterally symmetrical necrotic lesions in subcortical brain regions.
Defects in NDUFV1 are a cause of mitochondrial complex I deficiency (MT-C1D) [MIM:252010]. A disorder of the mitochondrial respiratory chain that causes a wide range of clinical disorders, from lethal neonatal disease to adult-onset neurodegenerative disorders. Phenotypes include macrocephaly with progressive leukodystrophy, non-specific encephalopathy, cardiomyopathy, myopathy, liver disease, Leigh syndrome, Leber hereditary optic neuropathy, and some forms of Parkinson disease.

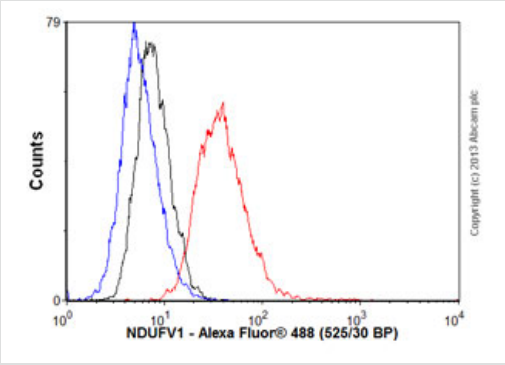
配列類似性

Belongs to the complex I 51 kDa subunit family.

細胞内局在

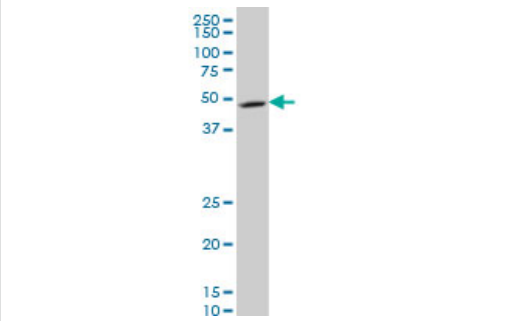
Mitochondrion inner membrane.

画像



Flow Cytometry - Anti-NDUFV1 antibody (ab55535)

Overlay histogram showing HepG2 cells stained with ab55535 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab55535, 0.1µg/1x10^6 cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-mouse IgG (H&L) (ab150113) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG2b [PLPV219] (ab91366, 1µg/1x10^6 cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter. This antibody gave a positive signal in HepG2 cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



Western blot - Anti-NDUFV1 antibody (ab55535)

NDUFV1 antibody (ab55535) at 1ug/lane + A-431 cell lysate at 25ug/lane.

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