abcam

Product datasheet

Anti-Ndufs4 antibody [EP7832] ab137064





重组 RabMAb

★★★★★ 13 Abreviews 6 References 14 图像

概述

产品名称 Anti-Ndufs4抗体[EP7832]

描述 兔单克隆抗体[EP7832] to Ndufs4

宿主 Rabbit

经测试应用 适用于: Flow Cyt (Intra), WB, IHC-P, ICC/IF, IP

种属反应性 与反应: Mouse, Rat, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

阳性对照 293T cell lysates, fetal brain and fetal stomach tissue lysates; Human brain and Human stomach

tissues Mouse heat lysate, rat heart lysate, mouse kidney, HeLa.

This product is a recombinant monoclonal antibody, which offers several advantages including: 常规说明

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb patents**.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

存储溶液 pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 0.05% BSA, 59% PBS

纯度 Protein A purified

单克降 克降 克隆编号 EP7832

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab137064于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Flow Cyt (Intra)		1/60. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody. For unpurified use at 1/100 - 1/1000 dilution.
WB	****(1)	1/500 - 1/10000. Detects a band of approximately 18 kDa (predicted molecular weight: 20 kDa).
IHC-P	★★★★★ (10)	1/50 - 1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
ICC/IF	★★★★ <u>(2)</u>	1/500. For unpurified use at 1/50 - 1/100.
IP		1/10 - 1/100.

靶	标

功能 Accessory subunit of the mitochondrial membrane respiratory chain NADH dehydrogenase

(Complex I), that is believed not to be involved in 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.

疾病相关 Defects in NDUFS4 are a cause of mitochondrial complex I deficiency (MT-C1D) [MIM:252010].

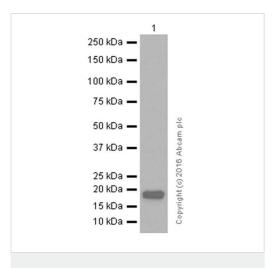
A disorder of the mitochondrial respiratory chain that causes a wide range of clinical disorders, from lethal neonatal disease to adult-onset neurodegenerative disorders. Phenotypes include macrocephaly with progressive leukodystrophy, non-specific encephalopathy, cardiomyopathy, myopathy, liver disease, Leigh syndrome, Leber hereditary optic neuropathy, and some forms of

Parkinson disease.

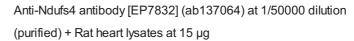
序列相似性 Belongs to the complex I NDUFS4 subunit family.

细**胞定位** Mitochondrion inner membrane.

图片



Western blot - Anti-Ndufs4 antibody [EP7832] (ab137064)

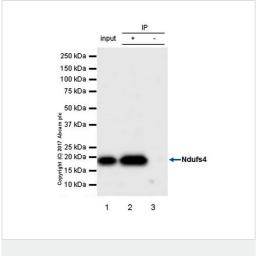


Secondary

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 20 kDa

Blocking and diluting buffer: 5% NFDM/TBST



Immunoprecipitation - Anti-Ndufs4 antibody [EP7832] (ab137064)

ab137064 (purified) at 1:30 dilution (2 μ g) immunoprecipitating Ndufs4 in Rat heart lysate.

Lane 1 (input): Rat heart lysate, 10µg

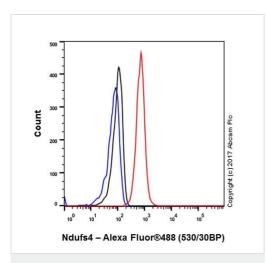
Lane 2 (+): ab137064 & Rat heart lysate

Lane 3 (-): Rabbit monoclonal IgG (ab172730) instead of

ab137064 in Rat heart lysate

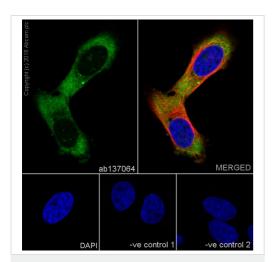
For western blotting, VeriBlot for IP Detection Reagent (HRP) (ab131366) was used for detection at 1:1000 dilution.

Blocking and diluting buffer: 5% NFDM/TBST.



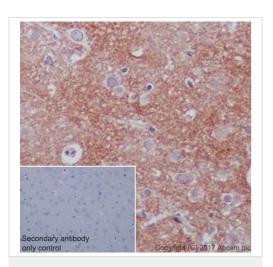
Flow Cytometry (Intracellular) - Anti-Ndufs4 antibody [EP7832] (ab137064)

Intracellular Flow Cytometry analysis of HeLa (Human cervix adenocarcinoma epithelial cell) cells labeling Ndufs4 with purified ab137064 at 1/60 dilution (red). Cells were fixed with 4% Paraformaldehyde. A Goat anti rabbit lgG (Alexa Fluor[®] 488) secondary antibody was used at 1/2000 dilution. Isotype control - Rabbit monoclonal lgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).



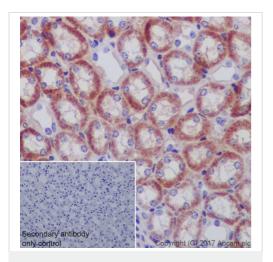
Immunocytochemistry/ Immunofluorescence - Anti-Ndufs4 antibody [EP7832] (ab137064)

Immunocytochemistry/ Immunofluorescence analysis of HeLa (Human cervix adenocarcinoma epithelial cell) cells labeling Ndusf5 with Purified ab137064 at 1:500 dilution. Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. Cells were counterstained with ab7291 anti-Tubulin (mouse mAb) ab150120 AlexaFluor ® 594 Goat anti-Mouse secondary (1:1000,2 µg/ml). ab150077 Goat anti rabbit lgG(Alexa Fluor ® 488) was used as the secondary antibody at 1:1000 dilution. DAPI nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



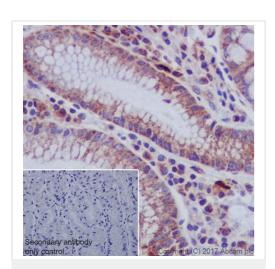
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ndufs4 antibody
[EP7832] (ab137064)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat cerebrum tissue sections labeling Ndufs4 with Purified ab137064 at 1:50 dilution. Heat mediated antigen retrieval was performed using Tris/EDTA buffer, pH 9.0. Tissue was counterstained with Hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.



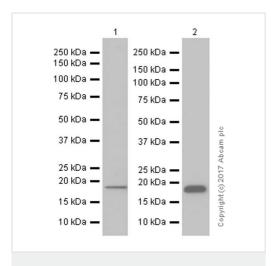
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ndufs4 antibody
[EP7832] (ab137064)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse kidney tissue sections labeling Ndufs4 with Purified ab137064 at 1:50 dilution. Heat mediated antigen retrieval was performed using Tris/EDTA buffer, pH 9.0. Tissue was counterstained with Hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ndufs4 antibody
[EP7832] (ab137064)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human stomach tissue sections labeling Ndufs4 with Purified ab137064 at 1:50 dilution. Heat mediated antigen retrieval was performed using Tris/EDTA buffer, pH 9.0. Tissue was counterstained with Hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.



Western blot - Anti-Ndufs4 antibody [EP7832] (ab137064)

All lanes : Anti-Ndufs4 antibody [EP7832] (ab137064) at 1/2000 dilution (purified)

Lane 1: Human fetal brain lysates

Lane 2: Mouse heart lysates

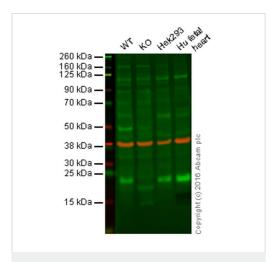
Lysates/proteins at 15 µg per lane.

Secondary

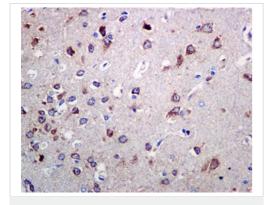
 $\begin{tabular}{ll} \textbf{All lanes:} Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution \end{tabular}$

Predicted band size: 20 kDa
Observed band size: 18 kDa

Blocking and diluting buffer: 5% NFDM/TBST.



Western blot - Anti-Ndufs4 antibody [EP7832] (ab137064)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ndufs4 antibody [EP7832] (ab137064)

Lane 1: Wild-type HAP1 cell lysate (20 µg)

Lane 2: Ndufs4 knockout HAP1 cell lysate (20 µg)

Lane 3: HEK293 cell lysate (20 µg)

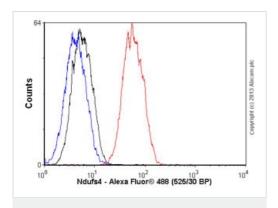
Lane 4: Human fetal heart tissue lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - ab137064 observed at 23 kDa. Red - loading control, ab8245, observed at 37 kDa.

Unpurified ab137064 was shown to recognize Ndufs4 when Ndufs4 knockout samples were used, along with additional cross-reactive bands. Wild-type and Ndufs4 knockout samples were subjected to SDS-PAGE. ab137064 and ab8245 (loading control to GAPDH) were diluted at 1/500 and 1/10 000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1/10000 dilution for 1 h at room temperature before imaging.

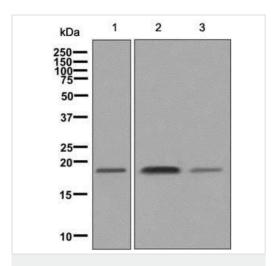
Immunohistochemical analysis of paraffin-embedded Human brain tissue labelling Ndufs4 with unpurified ab137064 at 1/50 dilution.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-Ndufs4 antibody [EP7832] (ab137064)

Overlay histogram showing HepG2 cells stained withunpurified ab137064 (red line). The cells were fixed with 80% methanol (5 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 (ab137064, 1/1000 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-rabbit lgG (H&L) (ab150077) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit IqG (monoclonal) (0.1µg/1x10⁶ 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 4% paraformaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



Western blot - Anti-Ndufs4 antibody [EP7832] (ab137064)

All lanes : Anti-Ndufs4 antibody [EP7832] (ab137064) at 1/1000 dilution (unpurified)

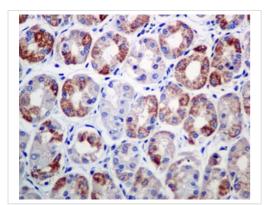
Lane 1: 293T cell lysate

Lane 2 : Fetal brain tissue lysate

Lane 3 : Fetal kidney tissue lysate

Lysates/proteins at 10 µg per lane.

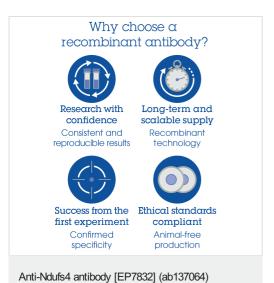
Predicted band size: 20 kDa **Observed band size:** 18 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ndufs4 antibody
[EP7832] (ab137064)

Immunohistochemical analysis of paraffin-embedded Human stomach tissue labelling Ndufs4 with unpurified ab137064 at 1/50 dilution.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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