# abcam

**Product datasheet** 

# Anti-NDUFA9 antibody [20C11B11B11] ab14713 敲除验证

★★★★★ 8 Abreviews 220 References 5 图像

## 概述

Anti-NDUFA9 <b>抗体</b> [20C11B11B11]		
小鼠单克隆抗体[20C11B11B11] to NDUFA9		
Mouse		
适用于: IHC-P, WB, Flow Cyt		
<b>与反</b> 应: Mouse, Rat, Cow, Human		
Tissue, cells or virus corresponding to Cow NDUFA9.		
WB: WI38 and NIH 3T3 whole cell lysates, human testis tissue lysate and human, cow, rat and mouse heart mitochondria. IHC-P: Human spinal column tissue. Flow Cyt: HepG2 cells.		
This monoclonal antibody to NDUFA9 has been knockout validated in Western blot. The expected band for NDUFA9 was observed in wild type cells and the band was not seen in knockout cells.		
This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or conjugation for your experiments, please contact <b>orders@abcam.com</b> .		
The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.		
If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As		

性能	
形式	Liquid
存 <b>放</b> 说明	Shipped at 4°C. Store at +4°C.
存储溶液	pH: 7.4 Preservative: 0.02% Sodium azide Constituent: HEPES buffered saline
纯 <b>度</b>	lgG fraction
纯化说明	Near homogeneity as judged by SDS-PAGE. The antibody was produced in vitro using
	1

	hybridomas grown in serum-free medium, and then purified by biochemical fractionation.
克隆	单 <b>克隆</b>
克 <b>隆</b> 编号	20C11B11B11
同种型	lgG1
轻链类 <b>型</b>	карра

应用

# The Abpromise guarantee

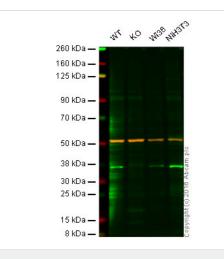
# Abpromise™承诺保证使用ab14713于以下的经测试应用

"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
IHC-P	★★★☆☆☆ <u>(2)</u>	Use at an assay dependent concentration.
WB	★ ★ ★ ★ ★ <u>(5)</u>	Use a concentration of 1 $\mu$ g/ml. Detects a band of approximately 36 kDa (predicted molecular weight: 40 kDa).
Flow Cyt		Use 1-2µg for 10 <sup>6</sup> cells. <u>ab170190</u> - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.

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.
序列相似性	Belongs to the complex I NDUFA9 subunit family.
细 <b>胞定位</b>	Mitochondrion matrix.

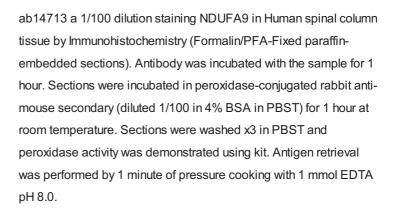
图片

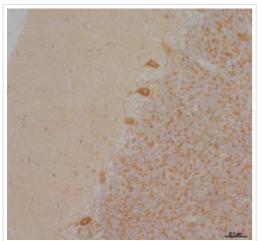


Western blot - Anti-NDUFA9 antibody [20C11B11B11] (ab14713)

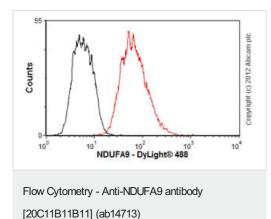
Lane 1: Wild-type HAP1 cell lysate (20 µg) Lane 2: NDUFA9 knockout HAP1 cell lysate (20 µg) Lane 3: WI38 cell lysate (20 µg) Lane 4: NIH3T3 cell lysate (20 µg) Lanes 1 - 4: Merged signal (red and green). Green - ab14713 observed at 40 kDa. Red - loading control, <u>ab176560</u>, observed at 52 kDa.

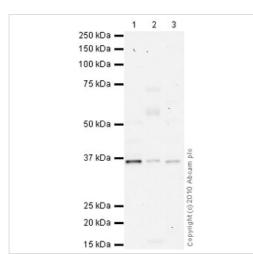
ab14713 was shown to specifically react with NDUFA9 in wild-type HAP1 cells. No band was observed when NDUFA9 knockout HAP1 samples were used. Wild-type and NDUFA9 knockout samples were subjected to SDS-PAGE. ab14713 and **ab176560** (loading control to alpha tubulin) were diluted at 1µg/mL and 1/10,000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Mouse IgG H&L (IRDye<sup>®</sup> 800CW) preadsorbed (**ab216772**) and Goat Anti-Rabbit IgG H&L (IRDye<sup>®</sup> 680RD) preadsorbed (**ab216777**) secondary antibodies at 1/10,000 dilution for 1 hour at room temperature before imaging.





Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NDUFA9 antibody [20C11B11B11] (ab14713)





Western blot - Anti-NDUFA9 antibody [20C11B11B11] (ab14713)

Overlay histogram showing HepG2 cells stained with ab14713 (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 (ab14713, 2 $\mu$ g/1x10<sup>6</sup> cells) for 30 min at 22°C. The secondary antibody used was DyLight<sup>®</sup> 488 goat anti-mouse IgG (H+L) (**ab96879**) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was Mouse IgG1 [ICIGG1] (**ab91353**, 2 $\mu$ g/1x10<sup>6</sup> cells) used under the same conditions. Acquisition of >5,000 events was performed.

All lanes : Anti-NDUFA9 antibody [20C11B11B11] (ab14713) at 1 µg/ml

Lane 1 : WI38 (Human lung fibroblast cell line) Whole Cell Lysate Lane 2 : Human testis tissue lysate - total protein (<u>ab30257</u>) Lane 3 : NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

## Secondary

**All lanes :** Goat polyclonal to Mouse lgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

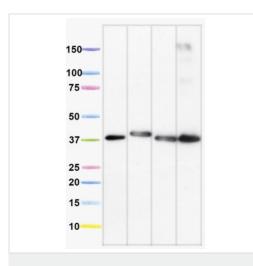
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 40 kDaObserved band size: 36 kDaAdditional bands at: 58 kDa. We are unsure as to the identity of these extra bands.

Exposure time: 20 minutes

The band observed at 36 kDa could potentially be a cleaved form of NDUFA9 due to the presence of a 35 amino acid transit peptide.



Western blot - Anti-NDUFA9 antibody [20C11B11B11] (ab14713) All lanes : Anti-NDUFA9 antibody [20C11B11B11] (ab14713)

Lane 1 : Isolated mitochondria from Human heart at 5  $\mu$ g Lane 2 : Isolated mitochondria from Bovine heart at 1  $\mu$ g Lane 3 : Isolated mitochondria from Rat heart at 10  $\mu$ g Lane 4 : Isolated mitochondria from Mouse Heart at 10  $\mu$ g

# Secondary

All lanes : Goat anti-Mouse IgG

Predicted band size: 40 kDa Observed band size: 37 kDa

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