# abcam

## Product datasheet

## Anti-NDUFB10 antibody [EPR16230-47] ab196019



重组 RabMAb

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概述

产品名称 Anti-NDUFB10抗体[EPR16230-47]

描述 兔单克隆抗体[EPR16230-47] to NDUFB10

宿主 Rabbit

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

种属反应性 与反应: Human

预测可用于: Mouse, Rat 🔷

免疫原 Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

阳性对照 HeLa, HepG2 and Jurkat cell lysates; Human transitional cell carcinoma of bladder tissue; HeLa

cells: HeLa whole cell extract.

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

- 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), 59% PBS, 0.05% BSA

纯度 Protein A purified

克隆 单克隆

克隆编号 EPR16230-47

**同种型** IgG

### 应用

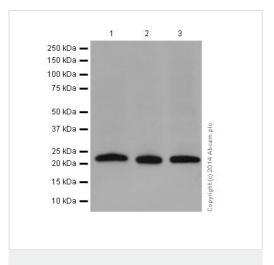
The Abpromise guarantee Abpromise™承诺保证使用ab196019于以下的经测试应用

"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
ICC/IF	**** <u>(1)</u>	1/350.
IP		1/50.
IHC-P		1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/10000. Detects a band of approximately 21 kDa (predicted molecular weight: 21 kDa).
Flow Cyt (Intra)		1/800.

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 NDUFB10 subunit family.
细胞定位	Mitochondrion inner membrane.

图片



Western blot - Anti-NDUFB10 antibody [EPR16230-47] (ab196019)

**All lanes :** Anti-NDUFB10 antibody [EPR16230-47] (ab196019) at 1/10000 dilution

**Lane 1 :** HeLa (Human epithelial cells from cervix adenocarcinoma) cell lysate

Lane 2 : HepG2 (Human liver hepatocellular carcinoma) cell lysateLane 3 : Jurkat (Human T cell leukemia cells from peripheral blood)cell lysate

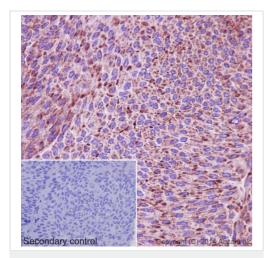
Lysates/proteins at 20 µg per lane.

#### Secondary

**All lanes :** Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 21 kDa **Observed band size:** 21 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

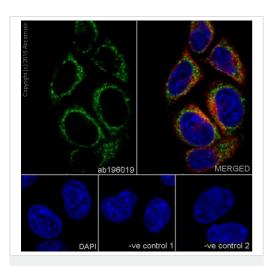


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NDUFB10 antibody
[EPR16230-47] (ab196019)

Immunohistochemical analysis of paraffin-embedded Human transitional cell carcinoma of bladder tissue labeling NDUFB10 with ab196019 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) secondary antibody at 1/500 dilution. Cytoplasm staining on Human transitional cell carcinoma of bladder tissue is observed. Counter stained with Hematoxylin.

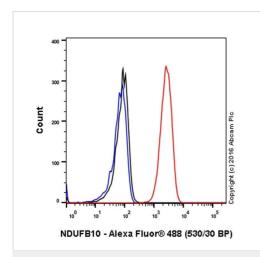
Secondary control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



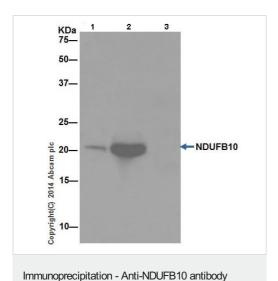
Immunocytochemistry/ Immunofluorescence - Anti-NDUFB10 antibody [EPR16230-47] (ab196019)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling NDUFB10 with ab196019 at 1/350 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Cytoplasm staining on HeLa cell line is observed. The nuclear counter stain is DAPI (blue).



Flow Cytometry (Intracellular) - Anti-NDUFB10 antibody [EPR16230-47] (ab196019)

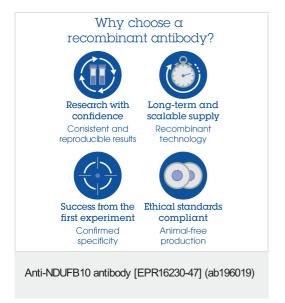
Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma) cells labelling NDUFB10 (red) with purified ab196019 at a dilution of 1/800. The secondary antibody used was Alexa Fluorr® 488 goat-anti-rabbit lgG (1/2000). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. Isotype control antibody was rabbit monoclonal lgG (black). The blue line shows cells without incubation with primary and secondary antibody.



[EPR16230-47] (ab196019)

NDUFB10 was immunoprecipitated from HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell extract with ab196019 at 1/50 dilution. Western blot was performed from the immunoprecipitate using ab196019 at 1/1000 dilution. Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG, was used as secondary antibody at 1/1500 dilution.

Lane 1: HeLa whole cell extract 10  $\mu$ g (Input). Lane 2: ab196019 IP in HeLa whole cell extract. Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab196019 in HeLa whole cell extract. Blocking and dilution buffer and concentration: 5% NFDM/TBST.



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