# abcam

# Product datasheet

# Anti-CENPF antibody ab5

★★★★★ 7 Abreviews 107 References 画像数 6

#### 製品の概要

製品名 Anti-CENPF antibody

製品の詳細 Rabbit polyclonal to CENPF

由来種 Rabbit

アプリケーション 適用あり: ICC, Flow Cyt, ICC/IF, IP, IHC-P, WB

種交差性 交差種: Mouse, Human

免疫原 Fusion protein with C-terminus of CENP-F (Human).

特記事項

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

#### 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** pH: 7.4

Preservative: 0.05% Sodium azide

Constituents: PBS, 50% Glycerol (glycerin, glycerine)

精製度 Immunogen affinity purified

**ポリ**/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab5の使用に適用されます

アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

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アプリケーション	Abreviews	特記事項
ICC	<b>★★★★ (1)</b>	Use at an assay dependent concentration.
Flow Cyt	**** (1)	Use at an assay dependent concentration. <b>ab171870</b> - Rabbit polyclonal lgG, is suitable for use as an isotype control with this antibody.
ICC/IF	<b>★★★★</b> ★ <u>(5)</u>	1/400 - 1/750.
IP		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration.
WB		1/1500. Detects a band of approximately 330 kDa (predicted molecular weight: 330 kDa). Block membranes for 1 hour with 5% nonfat dry milk/TBS-T.

## ターゲット情報

#### 機能

Required for kinetochore function and chromosome segregation in mitosis. Required for kinetochore localization of dynein, LIS1, NDE1 and NDEL1. Regulates recycling of the plasma membrane by acting as a link between recycling vesicles and the microtubule network though its association with STX4 and SNAP25. Acts as a potential inhibitor of pocket protein-mediated cellular processes during development by regulating the activity of RB proteins during cell division and proliferation. May play a regulatory or permissive role in the normal embryonic cardiomyocyte cell cycle and in promoting continued mitosis in transformed, abnormally dividing neonatal cardiomyocytes. Interaction with RB directs embryonic stem cells toward a cardiac lineage. Involved in the regulation of DNA synthesis and hence cell cycle progression, via its C-terminus. Has a potential role regulation skeletal myogenesis and in cell differentiation in embryogenesis. Involved in dendritic cell regulation of T-cell immunity against chlamydia.

関連疾患

配列類似性

発生段階

翻訳後修飾

細胞内局在

Stromme syndrome

Belongs to the centromere protein F family.

Gradually accumulates during the cell cycle, reaching peak levels in G2 and M phase, and is

rapidly degraded upon completion of mitosis.

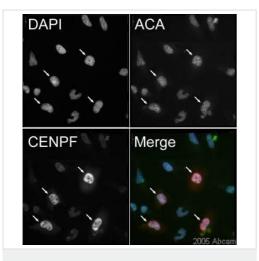
Hyperphosphorylated during mitosis.

Cytoplasm, perinuclear region. Nucleus matrix. Chromosome, centromere, kinetochore.

Cytoplasm, cytoskeleton, spindle. Relocalizes to the kinetochore/centromere (coronal surface of the outer plate) and the spindle during mitosis. Observed in nucleus during interphase but not in the nucleolus. At metaphase becomes localized to areas including kinetochore and mitotic apparatus as well as cytoplasm. By telophase, is concentrated within the intracellular bridge at

either side of the mid-body.

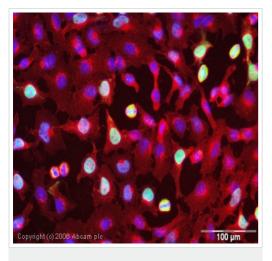
#### 画像



HeLa cells were labelled with anti-ACA and anti-CENPF (ab5). ab5 was used at a working dilution of 1/400. This image demonstrates the dramatic increase in fluorescence that occurs late in G2 cells (indicated by arrows). In the final panel DAPI is pseudo-coloured blue, while ACA and CENPF are coloured green and red respectively. 40x magnification.

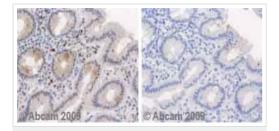
Immunocytochemistry/ Immunofluorescence - Anti-CENPF antibody (ab5)

This image is courtesy of Kirk McManus, University of British Columbia



Immunocytochemistry/ Immunofluorescence - Anti-CENPF antibody (ab5)

ICC/IF image of ab5 stained human HeLa cells. The cells were methanol fixed (5 min) and incubated with the antibody (ab5, 1µg/ml) for 1h at room temperature. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit lgG (H+L) used at a 1/1000 dilution for 1h. Image-iT<sup>TM</sup>FX Signal Enhancer was used as the primary blocking agent, 5% BSA (in TBS-T) was used for all other blocking steps. DAPI was used to stain the cell nuclei (blue). Alexa Fluor® 594 WGA was used to label plasma membranes (red).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CENPF antibody (ab5)

Ab5 staining Human normal colon tissue. Staining is localised to nuclear compartment.

Left panel: with primary antibody at 4 ug/ml. Right panel: isotype control.

Sections were stained using an automated system DAKO Autostainer Plus , at room temperature: sections were rehydrated and antigen retrieved with the Dako 3 in 1 AR buffer EDTA pH 9.0 in a DAKO PT Link. Slides were peroxidase blocked in 3% H2O2 in methanol for 10 mins. They were then blocked with Dako Protein block for 10 minutes (containing casein 0.25% in PBS) then incubated with primary antibody for 20 min and detected with Dako envision flex amplification kit for 30 minutes. Colorimetric detection

was completed with Diaminobenzidine for 5 minutes. Slides were counterstained with Haematoxylin and coverslipped under DePeX. Please note that for manual staining we recommend to optimize the primary antibody concentration and incubation time (overnight incubation), and amplification may be required.

DAPI
Interphase
Prophase

CENPF

Merge

Interpahse and Prophase HeLa cells were labelled with anti-ACA and anti-CENPF (ab5). ab5 was used at a working dilution of 1/400. This image emphasizes the redistribution of CENPF from the nuclear matrix during late G2 following entry into the initial stages of mitosis (see the accompanying image). Distinct punctate CENPF patterns proximally located in relation to the centromeres can be seen. In the final panel DAPI is pseudo-coloured blue, while ACA and CENPF are green and red respectively. 100x magnification.

Immunocytochemistry/ Immunofluorescence - Anti-CENPF antibody (ab5)

This image is courtesy of Kirk McManus, University of British Columbia

<u>kDa</u>
250 - **▼**150 - 100 - 100 - 50 - 25 -

Western blot - Anti-CENPF antibody (ab5)

All lanes: Anti-CENPF antibody (ab5) at 1/1500 dilution

Lane 1: Mitotic HeLa Lysate

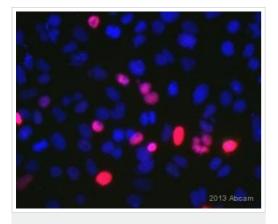
Lane 2 : Asynchronous HeLa Lysate

Lane 3: 50gamma Mitotic HeLa Lysate

Lane 4: 50gamma Asynchronous HeLa Lysate

Lysates/proteins at 25 µg per lane.

**Predicted band size:** 330 kDa **Observed band size:** 310 kDa



(human lung carcinoma cell line) cells stained for CENPF (red) using ab5 at 1/500 dilution in ICC/IF. ). Goat Anti-Rabbit lgG H&L (Alexa Fluor  $^{\!8}$  555) at 1/500 dilution was used as the secondary antibody.

Paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized A549

Immunocytochemistry/ Immunofluorescence - Anti-CENPF antibody (ab5)

This image is courtesy of an Abreview submitted by Dr. Stuart Rulten.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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