# abcam

## Product datasheet

## Anti-HCE antibody [EPR19384] ab201046



★★★★★ 5 Abreviews

10 Images

### Overview

**Product name** Anti-HCE antibody [EPR19384]

Rabbit monoclonal [EPR19384] to HCE **Description** 

**Host species** Rabbit

**Tested applications** Suitable for: IHC-P, WB, IP, ICC/IF Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Human fetal liver and fetal kidney lysates; Ramos, HepG2, SH-SY5Y, HeLa, C6, RAW 264.7,

> PC-12 and NIH/3T3 whole cell lysates; Mouse brain, kidney and spleen lysates; Rat brain and kidney lysates. IHC-P: Mouse cerebral cortex and rat spleen tissues. ICC/IF: HeLa and NIH/3T3

cells. IP: SH-SY5Y whole cell lysate.

General notes 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**.

### **Properties**

**Form** Liquid

Storage instructions 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.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

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

**Purity** Protein A purified

Clonality Monoclonal Clone number **EPR19384** 

**Isotype** IgG

## **Applications**

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab201046 in the following tested applications.

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

Application	Abreviews	Notes	
IHC-P		1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. The IHC application is recommended for mouse and rat only.	
WB	★★★★★ (5)	1/2000. Detects a band of approximately 69 kDa (predicted molecular weight: 69 kDa).	
IP		1/50.	
ICC/IF		1/500.	

I	aı	g	et

**Function** 

Bifunctional mRNA-capping enzyme exhibiting RNA 5'-triphosphatase activity in the N-terminal part and mRNA guanylytransferase activity in the C-terminal part. Catalyzes the first two steps of cap formation: by removing the gamma-phosphate from the 5'-triphosphate end of nascent mRNA to yield a diphosphate end, and by transferring the gmp moiety of GTP to the 5'-diphosphate terminus.

Tissue specificity

lsoform 1 and isoform 4 (at a lesser extent) are expressed in cerebrum, cerebellum, thyroid, lung,

heart, liver, kidney, spleen, large intestine, testis, skin and muscle.

Sequence similarities

In the N-terminal section; belongs to the non-receptor class of the protein-tyrosine phosphatase

family.

In the C-terminal section; belongs to the eukaryotic GTase family.

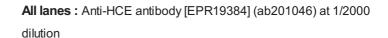
**Cellular localization** 

Nucleus.

## **Images**



Western blot - Anti-HCE antibody [EPR19384] (ab201046)



Lane 1: Human fetal liver lysate

Lane 2: Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

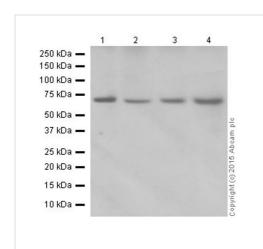
## Secondary

All lanes: Goat Anti-Rabbit IgG Peroxidase Conjugate, specific to the non-reduced form of IgG at 1/10000 dilution

Predicted band size: 69 kDa Observed band size: 69 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-HCE antibody [EPR19384] (ab201046)

All lanes: Anti-HCE antibody [EPR19384] (ab201046) at 1/2000 dilution

Lane 1: Ramos (Human Burkitt's lymphoma cell line) whole cell lysate

Lane 2: HepG2 (Human liver hepatocellular carcinoma cell line) whole cell lysate

Lane 3: SH-SY5Y (Human neuroblastoma cell line from bone marrow) whole cell lysate

Lane 4: HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

Lysates/proteins at 20 µg per lane.

## **Secondary**

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at

1/100000 dilution

Predicted band size: 69 kDa Observed band size: 69 kDa

Exposure time: 1 minute

Blocking/Dilution buffer: 5% NFDM/TBST.

250 kDa 250 kDa -150 kDa 🕳 150 kDa -100 kDa -100 kDa -75 kDa 🕳 75 kDa 🕳 50 kDa -50 kDa -37 kDa 🕳 37 kDa -25 kDa 🕳 25 kDa -20 kDa -20 kDa -15 kDa -15 kDa 🕳 10 kDa -10 kDa 🕳

Western blot - Anti-HCE antibody [EPR19384] (ab201046)

**All lanes :** Anti-HCE antibody [EPR19384] (ab201046) at 1/2000 dilution

Lane 1 : Mouse brain lysate

Lane 2: Mouse kidney lysate

Lane 3: Mouse spleen lysate

Lane 4 : Rat brain lysate

Lane 5: Rat kidney lysate

Lysates/proteins at 10 µg per lane.

## **Secondary**

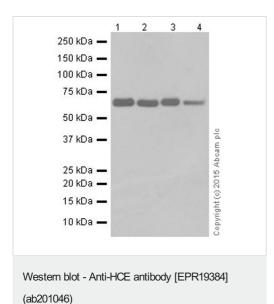
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at

1/100000 dilution

**Predicted band size:** 69 kDa **Observed band size:** 69 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



**All lanes :** Anti-HCE antibody [EPR19384] (ab201046) at 1/2000 dilution

Lane 1: C6 (Rat glial tumor cell line) whole cell lysate

Lane 2: RAW 264.7 (Mouse macrophage cell line transformed

with Abelson murine leukemia virus) whole cell lysate

**Lane 3 :** PC-12 (Rat adrenal gland pheochromocytoma cell line) whole cell lysate

Lane 4: NIH/3T3 (Mouse embryonic fibroblast cell line) whole cell lysate

Lysates/proteins at 10 µg per lane.

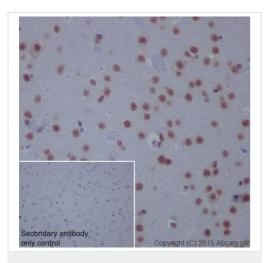
## **Secondary**

**All lanes :** Goat Anti-Rabbit  $\lg G \ H\&L \ (HRP) \ (\underline{ab97051})$  at 1/100000 dilution

Predicted band size: 69 kDa Observed band size: 69 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

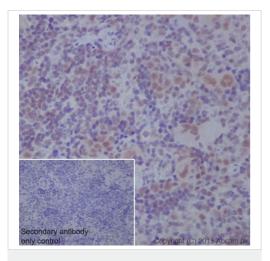


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HCE antibody
[EPR19384] (ab201046)

Immunohistochemical analysis of paraffin-embedded Mouse cerebral cortex tissue labeling HCE with ab201046 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nucleus staining on neurons of mouse cerebral cortex is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG 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.

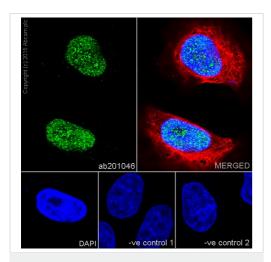


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HCE antibody
[EPR19384] (ab201046)

Immunohistochemical analysis of paraffin-embedded Rat spleen tissue labeling HCE with ab201046 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nucleus staining on lymphocytes of rat spleen is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG 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-HCE antibody [EPR19384] (ab201046)

ab201046 MERGED

DAPI -ve control 1 -ve control 2

Immunocytochemistry/ Immunofluorescence - Anti-HCE antibody [EPR19384] (ab201046)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling HCE with ab201046 at 1/500 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing nuclear staining on HeLa cell line. The nuclear counter stain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody [EPR19384] - Loading Control (ab7291) at 1/1000 dilution and Goat Anti-Mouse lgG H&L (Alexa Fluor® 594) preadsorbed (ab150120) at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: ab201046 at 1/500 dilution followed by **ab150120** at 1/1000 dilution.

-ve control 2: <u>ab7291</u> at 1/1000 dilution followed by <u>ab150077</u> at 1/1000 dilution.

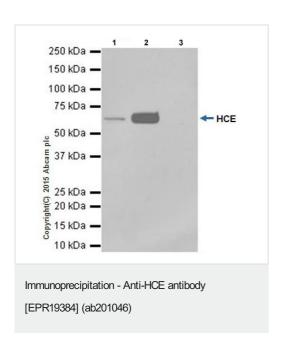
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized NIH/3T3 (Mouse embryonic fibroblast cell line) cells labeling HCE with ab201046 at 1/500 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing nuclear staining on NIH/3T3 cell line. The nuclear counter stain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody [EPR19384] - Loading Control (ab7291) at 1/1000 dilution and Goat Anti-Mouse lgG H&L (Alexa Fluor® 594) preadsorbed (ab150120) at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: ab201046 at 1/500 dilution followed by **ab150120** at 1/1000 dilution.

-ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> at 1/1000 dilution.



HCE was immunoprecipitated from 1mg of SH-SY5Y (Human neuroblastoma cell line from bone marrow) whole cell lysate with ab201046 at 1/50 dilution. Western blot was performed from the immunoprecipitate using ab201046 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/10000 dilution.

Lane 1: SH-SY5Y whole cell lysate 10µg (Input).

Lane 2: ab201046 IP in SH-SY5Y whole cell lysate.

Lane 3: Rabbit lgG,monoclonal [EPR19384] -lsotype Control (ab172730) instead of ab201046 in SH-SY5Y whole cell lysate. Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 30 seconds.



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

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