# abcam

# Product datasheet

# Anti-CSDE1/NRU antibody [EPR17413] ab200663

Recombinant RabMAb

1 References 8 Images

Overview

Product name Anti-CSDE1/NRU antibody [EPR17413]

**Description** Rabbit monoclonal [EPR17413] to CSDE1/NRU

Host species Rabbit

Tested applications Suitable for: WB, IP

Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: K562, HeLa, MCF7, C6, RAW 264.7 and PC-12 cell lysates; Human fetal brain lysate; HeLa

cytosolic fraction. IP: HeLa 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<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **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 EPR17413

**Isotype** IgG

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#### **Applications**

# The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab200663 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
WB		1/1000. Detects a band of approximately 89 kDa (predicted molecular weight: 89 kDa).
IP		1/140.

#### **Target**

**Function** RNA-binding protein. Required for internal initiation of translation of human rhinovirus RNA. May

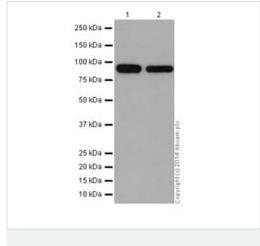
be involved in translationally coupled mRNA turnover. Implicated with other RNA-binding proteins in the cytoplasmic deadenylation/translational and decay interplay of the FOS mRNA mediated by

the major coding-region determinant of instability (mCRD) domain.

Sequence similarities Contains 9 CSD (cold-shock) domains.

Cellular localization Cytoplasm.

#### **Images**



Western blot - Anti-CSDE1/NRU antibody [EPR17413] (ab200663) **All lanes :** Anti-CSDE1/NRU antibody [EPR17413] (ab200663) at 1/2000 dilution

Lane 1: K562 (Human chronic myelogenous leukemia cells from

bone marrow) cell lysate

Lane 2: HeLa (Human epithelial cells from cervix

adenocarcinoma) cell lysate

Lysates/proteins at 20 µg per lane.

# **Secondary**

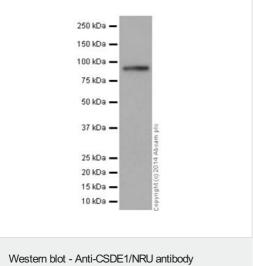
All lanes: Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at

1/1000 dilution

Predicted band size: 89 kDa Observed band size: 89 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Anti-CSDE1/NRU antibody [EPR17413] (ab200663) at 1/1000 dilution + MCF-7 (Human breast adenocarcinoma cell line) cell lysate at 20  $\mu g$ 

## **Secondary**

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

**Predicted band size:** 89 kDa **Observed band size:** 89 kDa

Exposure time: 3 minutes

[EPR17413] (ab200663)

Placking/Dilution buffer: 59/

Blocking/Dilution buffer: 5% NFDM/TBST.

250 kDa —

150 kDa —

100 kDa —

75 kDa —

37 kDa —

25 kDa —

20 kDa —

15 kDa —

10 kDa —

10 kDa —

Anti-CSDE1/NRU antibody [EPR17413] (ab200663) at 1/1000 dilution + Human fetal brain tissue lysate at 10 µg

# **Secondary**

Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

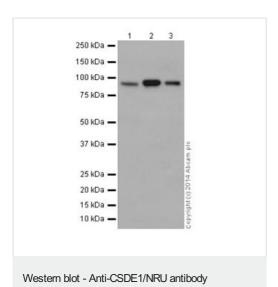
Predicted band size: 89 kDa Observed band size: 89 kDa

Exposure time: 3 minutes

[EPR17413] (ab200663)

Western blot - Anti-CSDE1/NRU antibody

Blocking/Dilution buffer: 5% NFDM/TBST.



[EPR17413] (ab200663)

**All lanes :** Anti-CSDE1/NRU antibody [EPR17413] (ab200663) at 1/1000 dilution

Lane 1: C6 (Rat glial tumor cells) cell lysate

Lane 2: RAW 264.7 (Mouse macrophage cells transformed with

Abelson murine leukemia virus) cell lysate

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

Lysates/proteins at 10 µg per lane.

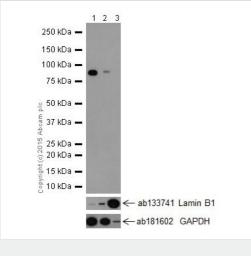
# **Secondary**

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

Predicted band size: 89 kDa Observed band size: 89 kDa

Exposure time: 1 minute

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-CSDE1/NRU antibody [EPR17413] (ab200663) **All lanes :** Anti-CSDE1/NRU antibody [EPR17413] (ab200663) at 1/2000 dilution

Lane 1: HeLa cytosolic fraction

Lane 2: HeLa membrane (mitochondrial) fraction

Lane 3: HeLa nuclear fraction

Lysates/proteins at 10 µg per lane.

## **Secondary**

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/50000 dilution

**Predicted band size:** 89 kDa **Observed band size:** 89 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

The HeLa cell fractions lysate set is ab168542 from Abcam.

1 2 3

250 kDa —

150 kDa —

100 kDa —

75 kDa —

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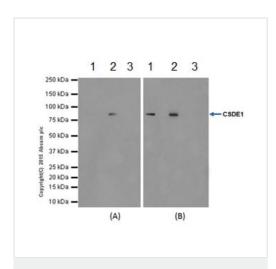
Immunoprecipitation - Anti-CSDE1/NRU antibody [EPR17413] (ab200663)

CSDE1/NRU was immunoprecipitated from 1mg of HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate with ab200663 at 1/140 dilution. Western blot was performed from the immunoprecipitate using ab200663 at 1/5000 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 lysate 10ug (Input). Lane 2: ab200663 IP in HeLa whole cell lysate. Lane 3: Rabbit monoclonal lgG (ab172730) instead of ab200663 in HeLa whole cell lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 30 seconds.



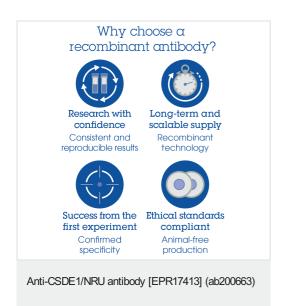
Immunoprecipitation - Anti-CSDE1/NRU antibody [EPR17413] (ab200663)

CSDE1/NRU was immunoprecipitated from 1mg of HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate with ab200663 at 1/140 dilution. Western blot was performed from the immunoprecipitate using ab200663 at 1/1000 dilution (Panel A) or ab201688 at 1/1000 dilution (Panel B). VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/1500 dilution.

Lane 1: HeLa whole cell lysate 10ug (Input). Lane 2: ab200663 IP in HeLa whole cell lysate. Lane 3: Rabbit monoclonal lgG (ab172730) instead of ab200663 in HeLa whole cell lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 3 min (Panel A); 10 seconds (Panel B).



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