


## Product datasheet

# Anti-CRISPR-Cas9 antibody [EPR19633] ab202657

Recombinant RabMAb

[1 References](#) [6 Images](#)

### Overview

<b>Product name</b>	Anti-CRISPR-Cas9 antibody [EPR19633]
<b>Description</b>	Rabbit monoclonal [EPR19633] to CRISPR-Cas9
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> WB, ICC/IF, IP, Flow Cyt (Intra)
<b>Species reactivity</b>	<b>Predicted to work with:</b> Streptococcus thermophilus 
<b>Immunogen</b>	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.
<b>Positive control</b>	WB: HEK-293 whole cell lysates transfected with CRISPR-Cas9 (G3ECR1, S.thermophilus, C-terminal aa801-1409) with GFP-Myc tag, CRISPR-Cas9 (G3ECR1) with Myc-His tag and CRISPR-Cas9 (Q03JL6, S.thermophilus (strain ATCC BAA-491 / LMD-9)) with Myc-His tag. ICC/IF: HeLa cells transfected with CRISPR-Cas9 (G3ECR1) with Myc-His tag. Flow Cyt: HEK-293 whole cell lysate transfected with CRISPR-Cas9 (G3ECR1) with Myc-His tag. IP: HEK-293 whole cell lysate transfected with CRISPR-Cas9 (G3ECR1) with Myc-His
<b>General notes</b>	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> <li>- High batch-to-batch consistency and reproducibility</li> <li>- Improved sensitivity and specificity</li> <li>- Long-term security of supply</li> <li>- Animal-free production</li> </ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a>.</p>

### Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	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.
<b>Storage buffer</b>	<p>pH: 7.2</p> <p>Preservative: 0.01% Sodium azide</p> <p>Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA</p>

<b>Purity</b>	Protein A purified
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	EPR19633
<b>Isotype</b>	IgG

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab202657 in the following tested applications.

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

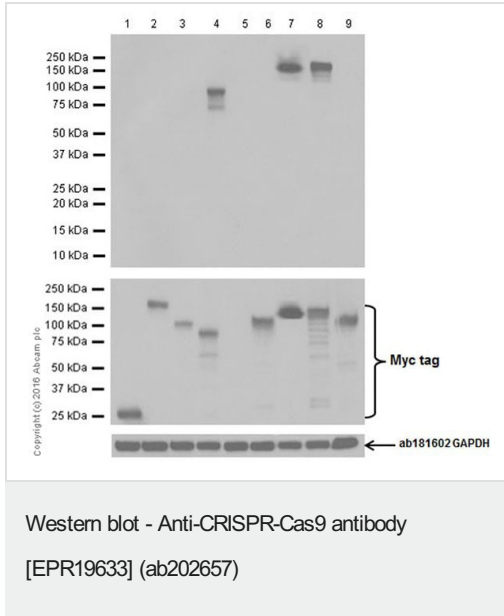
Application	Abreviews	Notes
<b>WB</b>		1/20000. Detects a band of approximately 164 kDa (predicted molecular weight: 164 kDa).
<b>ICC/IF</b>		1/100.
<b>IP</b>		1/30.
<b>Flow Cyt (Intra)</b>		1/60.

## Target

### Relevance

[FUNCTION] CRISPR (clustered regularly interspaced short palindromic repeat) is an adaptive immune system that provides protection against mobile genetic elements (viruses, transposable elements and conjugative plasmids). CRISPR clusters contain spacers, sequences complementary to antecedent mobile elements, and target invading nucleic acids. CRISPR clusters are transcribed and processed into CRISPR RNA (crRNA) (Probable). In type II CRISPR systems correct processing of pre-crRNA requires a trans-encoded small RNA (tracrRNA), endogenous ribonuclease 3 (rnc) and this protein. The tracrRNA serves as a guide for ribonuclease 3-aided processing of pre-crRNA. Subsequently Cas9/crRNA/tracrRNA endonucleolytically cleaves linear or circular dsDNA target complementary to the spacer. The target strand not complementary to crRNA is first cut endonucleolytically, then trimmed by 3'-5' exonucleolytically. DNA-binding requires protein and both RNA species. Cas9 probably recognizes a short motif in the CRISPR repeat sequences (the PAM or protospacer adjacent motif) to help distinguish self versus nonself.

## Images



**All lanes** : Anti-CRISPR-Cas9 antibody [EPR19633] (ab202657) at 1/20000 dilution

**Lane 1** : Empty vector with GFP-Myc tag (vector control) transfected HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate

**Lane 2** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (Q99ZW2, *Streptococcus pyogenes* serotype M1) with GFP-Myc tag

**Lane 3** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (G3ECR1, *Streptococcus thermophilus*, N-terminal aa1-800) with GFP-Myc tag

**Lane 4** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (G3ECR1, *Streptococcus thermophilus*, C-terminal aa801-1409) with GFP-Myc tag

**Lane 5** : Empty vector with Myc-His tag (vector control) transfected HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate

**Lane 6** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (A1IQ68, *Neisseria meningitidis* serogroup A / serotype 4A (strain Z2491)) with Myc-His tag

**Lane 7** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (G3ECR1, *Streptococcus thermophilus*) with Myc-His tag

**Lane 8** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (Q03J16, *Streptococcus thermophilus* (strain ATCC BAA-491 / LMD-9)) with Myc-His tag

**Lane 9** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (J7RUA5, *Staphylococcus aureus* subsp. *aureus*) with Myc-His tag

Lysates/proteins at 20 µg per lane.

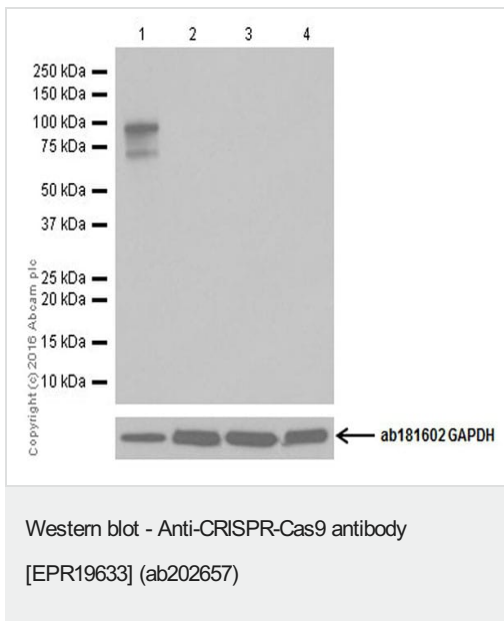
## Secondary

**All lanes** : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 100000 cells

**Predicted band size:** 164 kDa

**Exposure time:** 5 seconds

Blocking/Dilution buffer: 5% NFDm/TBST.



**All lanes** : Anti-CRISPR-Cas9 antibody [EPR19633] (ab202657) at 1/20000 dilution

**Lane 1** : HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (G3ECR1, Streptococcus thermophilus, C-terminal aa801-1409) with GFP-Myc tag

**Lane 2** : HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

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

**Lane 4** : Rat embryo lysate

Lysates/proteins at 20 µg per lane.

## Secondary

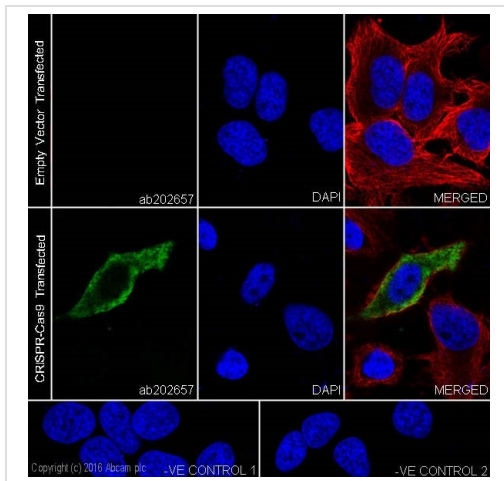
**All lanes** : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/100000 dilution

**Predicted band size:** 164 kDa

**Observed band size:** 86 kDa

**Exposure time:** 5 seconds

Blocking/Dilution buffer: 5% NFDm/TBST.



Immunocytochemistry/ Immunofluorescence - Anti-CRISPR-Cas9 antibody [EPR19633] (ab202657)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cell line from cervix adenocarcinoma) cells transfected with CRISPR-Cas9 (*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag or Empty vector, labeling CRISPR-Cas9 with ab202657 at 1/100 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor® 488) ([ab150079](#)) secondary antibody at 1/1000 dilution.

Confocal image showing Positive staining on HeLa cells transfected with CRISPR-Cas9(*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag.

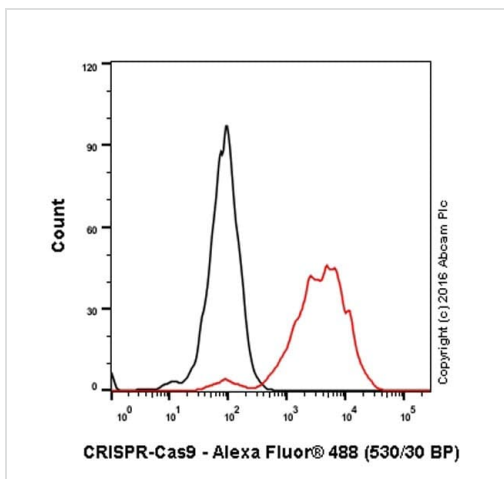
The nuclear counterstain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Loading Control ([ab7291](#)) at 1/1000 dilution and Goat Anti-Mouse IgG H&L (Alexa Fluor® 594) preadsorbed ([ab150120](#)) at 1/1000 dilution (red).

The negative controls are as follows:-

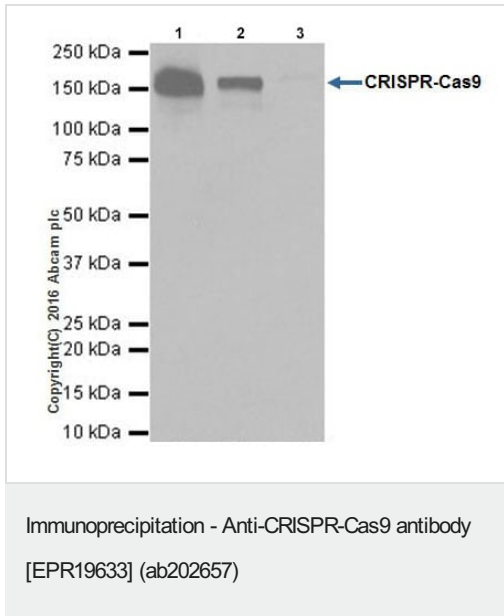
-ve control 1: ab202657 at 1/100 dilution followed by [ab150120](#) at 1/1000 dilution.

-ve control 2: [ab7291](#) at 1/1000 dilution followed by [ab150077](#) at 1/1000 dilution.



Flow Cytometry (Intracellular) - Anti-CRISPR-Cas9 antibody [EPR19633] (ab202657)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed HEK-293 (Human epithelial cell line from embryonic kidney) cells transfected with CRISPR-Cas9 (*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag labeling CRISPR-Cas9 with ab202657 at 1/60 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A]-Isotype control ([ab172730](#)) (black). Goat anti Rabbit IgG (Alexa Fluor® 488) at 1/2000 dilution was used as the secondary antibody.



CRISPR-Cas9 was immunoprecipitated from 0.35 mg of HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate transfected with CRISPR-Cas9 (*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag with ab202657 at 1/30 dilution.

Western blot was performed from the immunoprecipitate using ab202657 at 1/1000 dilution.

VeriBlot for IP Detection Reagent (HRP) (**ab131366**), was used for detection at 1/10000 dilution.

Lane 1: HEK-293 whole cell lysate transfected with CRISPR-Cas9 (*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag 10µg (Input).





Lane 2: ab202657 IP in HEK-293 whole cell lysate transfected with CRISPR-Cas9 (*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag.

Lane 3: Rabbit IgG, monoclonal [EPR25A]-Isotype Control (**ab172730**) instead of ab202657 in HEK-293 whole cell lysate transfected with CRISPR-Cas9 (*G3ECR1*, *Streptococcus thermophilus*) with Myc-His tag.

Blocking and dilution buffer and concentration: 5% NFD/MTBST.

Exposure time: 1 second.

Why choose a recombinant antibody?

 <p><b>Research with confidence</b> Consistent and reproducible results</p>	 <p><b>Long-term and scalable supply</b> Recombinant technology</p>
 <p><b>Success from the first experiment</b> Confirmed specificity</p>	 <p><b>Ethical standards compliant</b> Animal-free production</p>

Anti-CRISPR-Cas9 antibody [EPR19633] (ab202657)

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

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