

CRISPR/Cas9 Monoclonal Antibody

(Catalog No. A-9000)

Background:

The discovery of CRISPR (Clustered Regularly Interspaced Short Palindromic Repeats) and Cas9 (CRISPR associated system or CRISPR associated protein 9 nuclease) found in bacteria to work as a defense mechanism against foreign DNA, has proven to be an invaluable tool to target and modify a genetic sequence in gene editing and genome engineering applications. The system, known as CRISPR/Cas9, allows for sequence-specific cleavage of a targeted genomic locus by delivering the RNA-guided nuclease (Cas9) and appropriate guide RNAs (gRNA) into a cell. In addition, Protospacer Adjacent Motif (PAM) sequence immediately following the specificity sequence is necessary for successful binding of the Cas9 nuclease.

Concentration:

1 mg/ml

Description:

Mouse monoclonal antibody raised against CRISPR/Cas9, clone 7A9, generated with synthesized peptide corresponding to sequence of *Streptococcus pyogenes* (*S. pyogenes*) CRISPR-associated endonuclease Cas9/Csn1. This Anti-Cas9 mAb can detect CRISPR/Cas9 expression in target cells by WB, IF, IP, or ELISA to confirm and verify whether gRNA and Cas9 vectors are successfully transfected.

Specificity:

Recognizes both Cas9 and dCas9 (nuclease deficient Cas9)

Reactivity:

Species Independent

Isotype:

IgG1/Kappa

Formulation:

PBS, 30% Glycerol.

Storage:

Store at 4°C for short term (1-2 weeks). For long term storage, aliquot and then store at -20°C. Avoid multiple freeze/thaw cycles.

Purification:

Protein G purified

Application:

WB (1:1000), IP (2 µg/10⁶ cells), IF (1:500), ELISA

Ordering Information

Products:

CRISPR/Cas9 Monoclonal Antibody

Size

10 µg
50 µg
100 µg

Cat. No.

A-9000-010
A-9000-050
A-9000-100

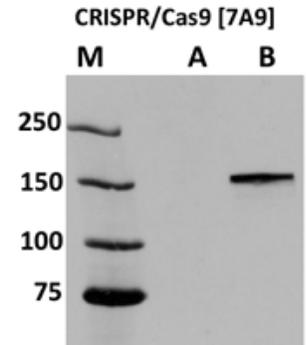


Fig1: Western Blot: Shown are the results of WB on protein extracts from transfected (B) and untransfected (A) HEK293 cells using the CRISPR/Cas9 [7A9] antibody.

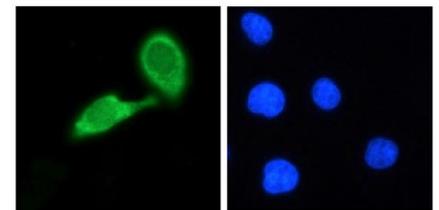


Fig2: Immunofluorescence: HeLa cells were transiently transfected with an N-terminally Flag-tagged *S. pyogenes* Cas9 expression vector. The cells were stained with the Cas9 antibody followed by anti mouse-AF488 coupled secondary antibody. Nuclei were counter-stained with Hoechst 33342.

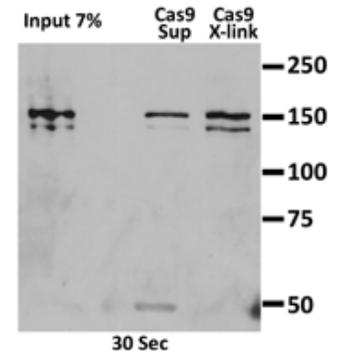


Fig3 Immunoprecipitation: HEK293T expressing N-terminally Flag-tagged *S.pyogenes* Cas9 were lysed 72H post transfection. Proteins were immunoprecipitated from 100 µg of whole cell lysate for 1h at 4°C with Cas9 Sup followed by incubation for 1h at 4°C with a 1:1 mixture of protein A/G sepharose beads, or for 2h at 4°C with Cas9 ab X-linked to a 1:1 mixture of protein A/G sepharose beads. Beads were washed and eluted by boiling in Laemmli, separated by SDS-PAGE, and transferred to nitrocellulose. Membrane was blocked, incubated with Cas9 ab, then incubated with HRP anti-mouse antibody.

This product is for research purposes only. Not intended for use in diagnostic procedures.