

MODIFICATIONS TO CAS₉ FOR PASSIVE-DELIVERY INTO CELLS

Tech ID: 25286 / UC Case 2016-016-0

PATENT STATUS

Country	Type	Number	Dated	Case
United States Of America	Issued Patent	11,118,194	09/14/2021	2016-016

BRIEF DESCRIPTION

RNA-mediated adaptive immune systems in bacteria and archaea rely on Clustered Regularly Interspaced Short Palindromic Repeat (CRISPR) genomic loci and CRISPR-associated (Cas) proteins that function together to provide protection from invading viruses and plasmids. In Type II CRISPR-Cas systems, the Cas9 protein functions as an RNA-guided endonuclease that uses a dual-guide RNA consisting of crRNA and trans-activating crRNA (tracrRNA) for target recognition and cleavage by a mechanism involving two nuclease active sites that together generate double-stranded DNA breaks (DSBs). Thus, the Cas9 system provides a facile means of modifying genomic information.

UC Berkeley researchers have developed modified site-directed modifying polypeptides and ribonucleoproteins comprising the modified polypeptides. As the modified site-directed modifying polypeptides are modified for passive entry into target cells, the polypeptides are useful in a variety of methods for target nucleic acid modification.

SUGGESTED USES

- » Genome editing
- » Gene therapy

ADVANTAGES

- » Crosses the plasma membrane of a eukaryotic cell without the need for any additional agent (e.g., small molecule agents, lipids, etc.)

ADDITIONAL TECHNOLOGIES BY THESE INVENTORS

- COMPOSITIONS AND METHODS FOR IDENTIFYING HOST CELL TARGET PROTEINS FOR TREATING RNA VIRUS INFECTIONS
- Genome Editing via LNP-Based Delivery of Efficient and Stable CRISPR-Cas Editors
- Type III CRISPR-Cas System for Robust RNA Knockdown and Imaging in Eukaryotes
- Cas12-mediated DNA Detection Reporter Molecules
- Improved guide RNA and Protein Design for CasX-based Gene Editing Platform
- Cas13a/C2c2 - A Dual Function Programmable RNA Endoribonuclease
- RNA-directed Cleavage and Modification of DNA using CasY (CRISPR-CasY)
- CasX Nickase Designs, Tans Cleavage Designs & Structure

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INVENTORS

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OTHER INFORMATION

KEYWORDS

CRISPR, Cas9, gene, genome editing

CATEGORIZED AS

- » **Biotechnology**
- » Genomics
- » **Medical**
- » Gene Therapy
- » Research Tools
- » **Research Tools**
- » Nucleic Acids/DNA/RNA

RELATED CASES

2016-016-0

- ▶ [In Vivo Gene Editing Of Tau Locus Via Liponanoparticle Delivery](#)
- ▶ [A Dual-RNA Guided CasZ Gene Editing Technology](#)
- ▶ [CRISPR-CAS EFFECTOR POLYPEPTIDES AND METHODS OF USE THEREOF \(“Cas-VariPhi”\)](#)
- ▶ [A Protein Inhibitor Of Cas9](#)
- ▶ [RNA-directed Cleavage and Modification of DNA using CasX \(CRISPR-CasX\)](#)
- ▶ [Compositions and Methods for Genome Editing](#)
- ▶ [Split-Cas9 For Regulatable Genome Engineering](#)
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- ▶ [Methods to Interfere with Prokaryotic and Phage Translation and Noncoding RNA](#)
- ▶ [CRISPR CASY COMPOSITIONS AND METHODS OF USE](#)
- ▶ [Single Conjugative Vector for Genome Editing by RNA-guided Transposition](#)
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- ▶ [Engineered/Variant Hyperactive CRISPR CasPhi Enzymes And Methods Of Use Thereof](#)
- ▶ [Engineering Cas12a Genome Editors with Minimized Trans-Activity](#)
- ▶ [Methods Of Use Of Cas12L/CasLambda In Plants](#)
- ▶ [Type V CRISPR/CAS Effector Proteins for Cleaving ssDNA and Detecting Target DNA](#)
- ▶ [THERMOSTABLE RNA-GUIDED ENDONUCLEASES AND METHODS OF USE THEREOF \(GeoCas9\)](#)
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- ▶ [Endoribonucleases For Rna Detection And Analysis](#)
- ▶ [Efficient Site-Specific Integration Of New Genetic Information Into Human Cells](#)
- ▶ [CRISPR-Cas Effector Polypeptides and Methods of Use Thereof](#)
- ▶ [Class 2 CRISPR/Cas COMPOSITIONS AND METHODS OF USE](#)
- ▶ [Compositions and Methods of Use for Variant Csy4 Endoribonucleases](#)
- ▶ [Identification Of Sites For Internal Insertions Into Cas9](#)
- ▶ [Methods and Compositions for Controlling Gene Expression by RNA Processing](#)



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