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Highly efficient base editing in *Staphylococcus aureus* using an engineered CRISPR RNA-guided cytidine deaminase†

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Novel therapeutic means against *Staphylococcus aureus* infections are urgently needed due to the emergence of drug-resistant *S. aureus*. We report the development of a CRISPR RNA-guided cytidine deaminase (pnCasSA-BEC), enabling highly efficient gene inactivation and point mutations in *S. aureus*. We engineered a fusion of a Cas9 nickase (Cas9D10A) and a cytidine deaminase (APOBEC1) that can be guided to a target genomic locus for gene inactivation *via* generating a premature stop codon. The pnCasSA-BEC system nicks the non-edited strand of the genomic DNA, directly catalyzes the conversion of cytidine (C) to uridine (U), and relies on DNA replication to achieve C → T (G → A) conversion without using donor repair templates. The development of the base-editing system will dramatically accelerate drug-target exploration in *S. aureus* and provides critical insights into the development of base-editing tools in other microbes.

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Introduction

As a major human pathogen, *Staphylococcus aureus* is the leading cause of hospital- and community-acquired infections. This pathogen can cause a wide variety of infections, including minor skin infections and life-threatening diseases, such as endocarditis, necrotizing pneumonia, and toxic shock syndrome.^{1,2} The emergence of drug-resistant *S. aureus*, such as methicillin-resistant and vancomycin-intermediate *S. aureus* (MRSA and VISA), has posed a severe public crisis worldwide.^{3,4} Hence it is urgently needed to develop novel therapeutic means against drug-resistant *S. aureus* infections.

We have developed a CRISPR/Cas9-mediated genome editing system (pCasSA).⁵ This method allows for rapid and efficient genetic manipulation in *S. aureus*, accelerating bacterial physiology studies, such as pathogenesis and drug resistance, and boosting novel drug-target exploration and new therapeutic method development.

The CRISPR/Cas9 system was originally discovered in bacterial immune systems and has been engineered for genome editing in a variety of organisms.^{5–15} The system is composed of a Cas9 nuclease and a synthetic guide RNA (sgRNA) (Fig. 1A). The Cas9 nuclease forms a complex with sgRNA and can be

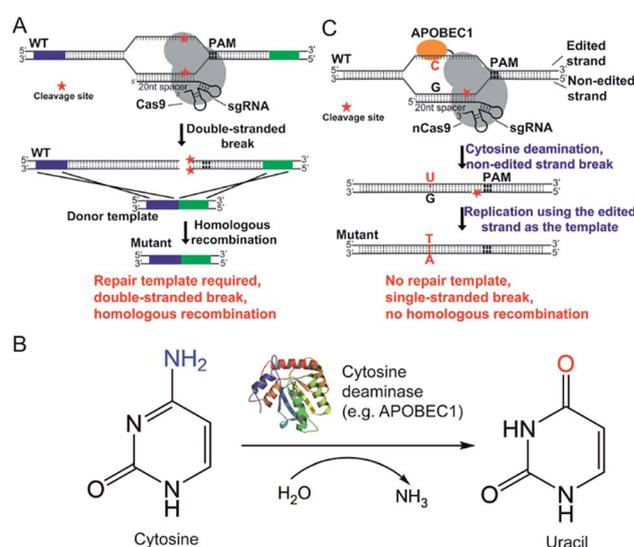


Fig. 1 Comparison between the CRISPR/Cas9-mediated genome editing method and the “cytidine base editor”-mediated genome editing method. (A) Detailed mechanism of the CRISPR/Cas9-mediated genome editing method. (B) The deamination reaction catalyzed by the cytosine deaminase. (C) Detailed mechanism of the “base editor” APOBEC1-nCas9-mediated genome editing method.

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directed to any genomic locus *via* base pairing of the 20-nucleotide sequence at the 5' prime of sgRNA with the genomic DNA when an adjacent protospacer-adjacent motif (PAM) is present in the target locus (Fig. 1A). After targeting, the Cas9 nuclease cleaves the genomic DNA by its enzymatic active residues (Asp10 and His840 in the case of *Streptococcus pyogenes* Cas9), generating a double-stranded break in the target genome (Fig. 1A). Given that the double-stranded DNA break is lethal to bacteria because of the lack of the non-homologous end-joining (NHEJ) DNA repair mechanism, only the cells that undergo the homologous recombination repair process can survive after editing. Hence the pCasSA system is capable of selecting separate double-crossover events that occur in the traditional *S. aureus* genome editing methods (e.g. pKOR1, pMAD, and pIMAY)^{16–18} in one step, achieving convenient genome editing in *S. aureus* (Fig. 1A). However, the weak intrinsic homologous recombination capacity of *S. aureus* prevents the high-rate recovery of survival cells after the double-stranded break of the genome, thus hampering the applications of the pCasSA system in strains with low transformation efficiencies, such as many MRSA strains directly isolated from patients.

Fortunately, the recent development of “base editors” allows for direct mutagenesis of bases inside the genome, opening a new avenue for genome editing in biology.^{19–24} Until now, two kinds of base editors have been developed (the cytidine editor and the adenosine editor) and each one is composed of a dead Cas9 (Cas9D10A or H840A) protein or a Cas9 nickase (Cas9D10A) and a deaminase fused to the Cas9 protein (Fig. 1B and C). The deaminase can be directed to any genomic locus by the Cas9/sgRNA complex to achieve base editing in the single-stranded DNA generated upon Cas9/sgRNA binding (Fig. 1B and C). The CRISPR/Cas9 genome editing method creates a double-stranded break, relies on the homologous recombination repair mechanism (in bacteria), and requires a donor repair template. In contrast, base editors directly catalyze the conversion of cytidine to uridine through a deamination reaction (adenosine to guanosine conversion in the case of recently developed adenine editors), nick the non-edited strand, and use the DNA replication mechanism to achieve editing (Fig. 1B and C). Hence we envision that “base editors” will allow for efficient genetic manipulation in *S. aureus* without sacrificing transformation CFUs.

In this study, we have developed a highly efficient and convenient base-editing system (pnCasSA–BEC) in *S. aureus* *via* engineering a fusion of a Cas9 nickase and a cytidine deaminase. The tool allows for gene inactivation and point mutations in the genomes of *S. aureus*, thereby dramatically accelerating bacterial physiology studies and drug-target exploration. The development of the pnCasSA–BEC system will provide critical insights into base-editing system development in other microbes.

Results and discussion

To harness “base editors” for base editing in *S. aureus*, we designed and constructed the plasmid pnCasSA–BEC (Fig. 2A). Inspired by the success of using *cap 1A* and *rpsL* promoters to drive the expressions of the sgRNA and the Cas9 protein,

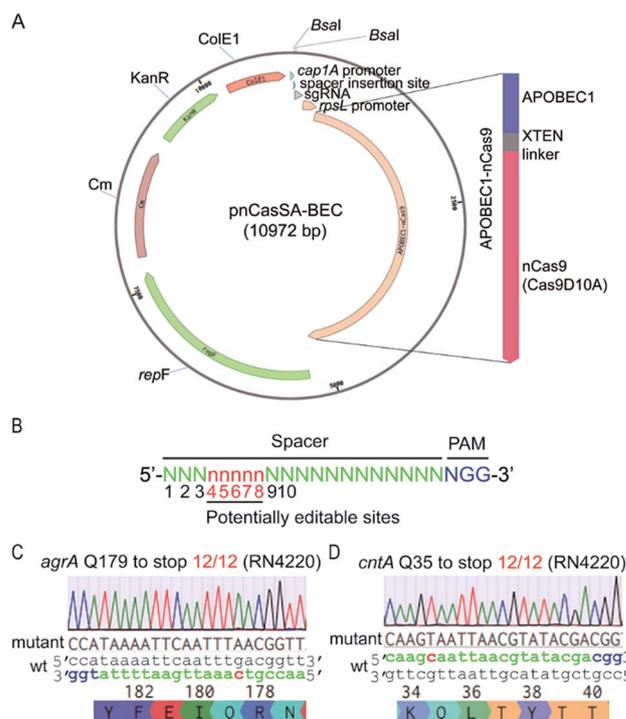


Fig. 2 The engineered base-editing system enables highly efficient C → T conversion in the *S. aureus* RN4220 strain. (A) Map of the base-editing plasmid pnCasSA–BEC. APOBEC1–nCas9, a fusion protein composed of a deaminase APOBEC1 at the N terminus and a Cas9 nickase Cas9D10A at the C terminus; *rpsL* promoter, an *S. aureus* strong promoter used to drive the expression of APOBEC1–nCas9; *BsaI* sites, Golden Gate assembly of spacers; *cap 1A* promoter, the sgRNA expression promoter; *ColE1*, an *E. coli* replication origin. KanR, the antibiotic marker kanamycin used in *E. coli*; Cm, the chloramphenicol-resistance marker used in *S. aureus*; *repF*, an *S. aureus* temperature-sensitive origin for plasmid curing after editing. (B) The potential editable sites in the genome. The prediction of the potential editable sites is based on the mammalian “base editor” BE3.¹⁹ (C and D) The pnCasSA–BEC system enables highly efficient base editing in the RN4

The mammalian C → T conversion “base editor” has an editable window from positions 4 to 8 in the spacer sequence (Fig. 2B) with the favorable editing site at position 7.¹⁹ To assess the capacity of the pnCasSA-BEC system in the conversion of C → T in *S. aureus*, we assembled different spacers of *agrA*, *cntA*, and *esaD* genes that contain potentially editable C(s). Next, we transformed the assembled plasmids into the laboratory *S. aureus* strain RN4220. As shown in Fig. 2C, D and S1,† the C at position 7 of *agrA*, the C at position 5 of *cntA*, and the C at position 6 of *esaD* were mutated to T successfully with 100% efficiency, resulting in the generation of premature stop codons. However, the Cs at positions 3 and 4 of *agrA* (Fig. 2C) and the C at position 3 of *esaD* (Fig. S1†) could not be mutated to T in the experiment.

Because of the different capacities of the pnCasSA-BEC system in mutating C(s) at different positions, we systematically investigated the editing window of the pnCasSA-BEC system in *S. aureus*. We assembled 11 different spacers containing C(s) at different positions (from positions 2 to 12) into the pnCasSA-BEC plasmid and transformed them individually into the RN4220 strain to examine the editing efficiencies. As shown in Fig. S2,† the Cs at positions 4 to 8 were mutated to Ts with 100% efficiency whereas the Cs at positions 2, 3, and 9–12 could not be mutated to T by the pnCasSA-BEC system, indicating that the editable window of the pnCasSA-BEC system is from positions 4 to 8. The discrepancy in the editing efficiencies at position 4 of the *agrA* spacer (Fig. 2C) and spacer 3 (Fig. S2†) indicates that other factors, such as adjacent bases or the sequence content of the spacers, can affect the editing efficiency of the pnCasSA-BEC system at position 4. In fact, the *in vitro* activity of the base editors follows the order TC ≥ CC ≥ AC > GC.¹⁹

Achieving a desired amino acid mutation may require the mutations of multiple Cs. We thus assessed the capacity of the pnCasSA-BEC system in mutating two successive Cs in *S. aureus*. Four different spacers containing two successive Cs at positions 4–5, 5–6, 6–7, and 7–8 were assembled into the pnCasSA-BEC plasmid, respectively. They were transformed into the RN4220 strain to test the editing efficiencies. As shown in Fig. S3,† both the successive Cs except the C at position 4 of spacer 1 could be mutated to Ts by the pnCasSA-BEC system with high efficiencies from 4/6 to 6/6 (or 18/18), demonstrating the great capacity of the pnCasSA-BEC system in mutating two successive Cs in the editable window. The C at position 4 of spacer 1 could not be mutated to T by the pnCasSA-BEC system (Fig. S3†).

The pnCasSA-BEC plasmid contains the same temperature-sensitive replication origin *repF* as the pCasSA plasmid. We used the same method⁵ to cure the pnCasSA-BEC plasmid after editing as that of the pCasSA plasmid. As shown in Fig. S4,† after culturing the cells at a non-permissive temperature (42 °C), all four randomly picked colonies could grow normally in the absence of chloramphenicol whereas none of them could grow in the presence of the antibiotic, confirming the successful removal of the pnCasSA-BEC plasmid from the cells (Fig. S4A and B†).

In agree with our expectation, the transformation of the spacer-introduced pnCasSA-BEC plasmid did not reduce the CFU (Fig. S5A and B†). However, the transformation of the

spacer- or both the spacer and repair arm-introduced pCasSA plasmid drastically reduced the CFU (Fig S5A and B†).⁵ Thereby the pnCasSA-BEC system may possess a greater capacity than the pCasSA system in genome editing in strains with low transformation efficiencies, such as some MRSA strains directly isolated from patients.

Next, we tested the base-editing efficiency of the pnCasSA-BEC system in four clinically isolated *S. aureus* strains, including ST239 (770), ST398 (1059), Newman, and USA300. ST239 (770) and ST398 (1059) are two MRSA strains isolated from patients in Ren Ji Hospital, China.^{27,28} The *agrA*-spacer and the *cntA*-spacer introduced pnCasSA-BEC plasmids isolated from the RN4220 strain were transformed into the four aforementioned strains by electroporation.²⁹ Several *S. aureus* restriction-system-modified *E. coli* strains, such as IM08B and ALC7885, probably can also be used for the restriction modification of the pnCasSA-BEC plasmid.^{30,31} As shown in Fig. 3A–D and S6A–D,† the efficiencies of C → T conversions at position 7 of the *agrA* spacer and position 5 of the *cntA* spacer in all four strains were 100%, demonstrating the great capacity of the pnCasSA-BEC system in base editing in the clinically isolated *S. aureus* strains. Because the disruption of the *agrA* gene in *S. aureus* abolishes the pathogen's hemolysis activity,³² the C → T mutation in the *agrA* gene (generation of a premature stop codon) in all four strains was also confirmed phenotypically by the hemolysis assay (Fig. 3A, C and S6A, C†). In agree with the results obtained in the RN4220 strain, the spacer-introduced

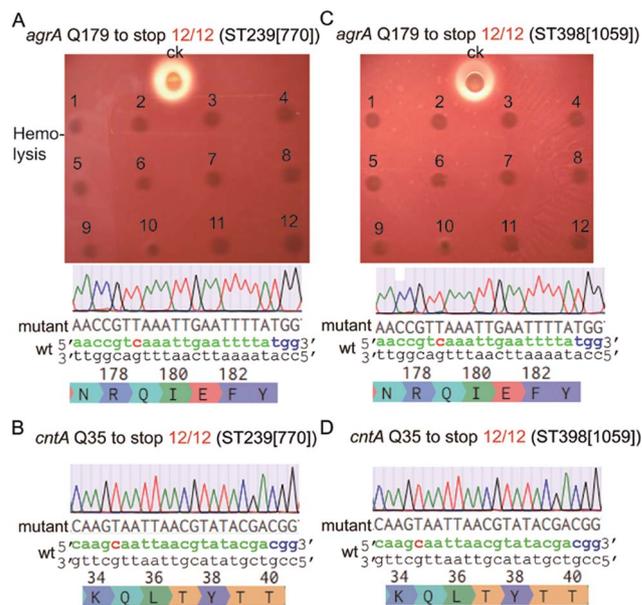


Fig. 3 The pnCasSA-BEC system enables highly efficient C → T conversion in the clinically isolated *S. aureus* strains. (A and B) Q179 of the *agrA* gene (A) and Q35 of the *cntA* gene (B) of the *S. aureus* ST239 (770) strain were successfully mutated to stop codons with 100% efficiency by using the pnCasSA system. ck: the wild-type strain of ST239 (770). (C and D) Q179 of the *agrA* gene (C) and Q35 of the *cntA* gene (D) of the *S. aureus* ST398 (1059) strain were successfully mutated to stop codons with 100% efficiency by using the pnCasSA system. ck: the wild-type strain of ST398 (1059).



pnCasSA-BEC plasmid exhibited similar transformation efficiencies to the empty pnCasSA-BEC plasmid in all four strains (Fig S7A and B†).

To gain insights into overall editable sites of the pnCasSA-BEC system in *S. aureus*, we performed bioinformatics scanning against the genomes of the MRSA252 and Newman strains. The results showed that almost all the genes (98.81% of MRSA252 and 97.6% of Newman) contain at least one PAM site (Fig. 4A). 96.12% of the genes of the MRSA252 strain and 95.7% of the genes of the Newman strain contain possibly editable C(s)

(Fig. 4A). The potential codons that can be changed and the potential changed amino acids are listed in Fig. S8A and B.† In addition, 68.8% of the genes of the MRSA252 strain and 70.36% of the genes of the Newman strain possess possibly editable stop sites (CAA [Gln], CAG [Gln], CGA [Arg], and TGG [Trp]) (Fig. 4A). All the possibly editable stop sites are listed in Tables S1 and S2.† Because the closer the editable stop sites are to start codons, the higher possibility the translated product would be inactive, we analyzed the distribution of the possibly editable stop sites inside the genes. As shown in Fig. 4B, in the Newman strain, 29%, 52.55%, and 64.8% of the possibly editable stop sites locate at the top 20%, 50%, and 80% of the gene bodies, respectively, while in the MRSA252 strain, 29.2%, 51.6%, and 63.5% of the possibly editable stop sites locate at the top 20%, 50%, and 80% of the gene bodies, respectively. Further bioinformatics analysis revealed that the editable sites of the pnCasSA-BEC system are almost evenly distributed across the genome without chain preference (Fig. 4C). Together, these analyses unveiled that numerous editable sites of the pnCasSA-BEC system are present in *S. aureus* genomes and many genes can be inactivated using this system.

Conclusions

We have developed a highly efficient and convenient base-editing system in *S. aureus* by engineering a fusion of a Cas9 nickase and a cytidine deaminase. Unlike our previously developed pCasSA system, the pnCasSA-BEC system can achieve efficient genome editing without using repair templates or sacrificing transformation CFUs. Given its ease of use and high efficiency, this system can probably be further engineered to be a multiplexed editing system that cannot be achieved by the pCasSA system. Besides gene inactivation, the pnCasSA-BEC system is also capable of editing other sites in the genomes, such as the promoter regions and the regions coding for small RNAs. Future applications of the pnCasSA-BEC system can dramatically accelerate both the basic-science and applied investigations in *S. aureus*, in particular in strains with low-transformation efficiencies. The development of the pnCasSA-BEC system will shed light on base-editing system development in other microbes.

Experimental

Strains, plasmids, primers, and growth conditions

Plasmids and strains used in this study are listed in Table S3.† Primers used in this study are listed in Table S4.† *E. coli* strains and *S. aureus* strains were grown in Luria-Bertani broth (LB) and tryptic soy broth (TSB), respectively. Antibiotics were used at the following concentrations: kanamycin 50 $\mu\text{g mL}^{-1}$ for *E. coli*, chloramphenicol 5 $\mu\text{g mL}^{-1}$ for the *S. aureus* RN4220 strain, and 10 $\mu\text{g mL}^{-1}$ for other *S. aureus* strains.

Construction of the pnCasSA-BEC plasmid

To achieve the construction of the pnCasSA-BEC plasmid, we first constructed the pnCasSA plasmid, which contained a Cas9

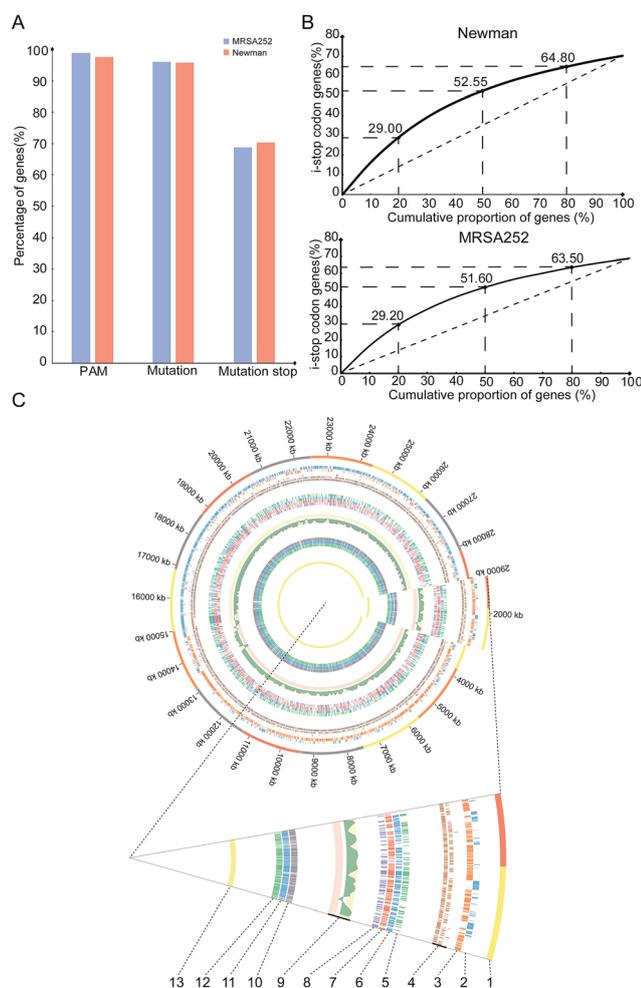


Fig. 4 Predicted editable sites of the pnCasSA-BEC system across the entire genome of *S. aureus*. (A) Proportion of genes containing predicted editable sites of the pnCasSA-BEC system. PAM: PAM sites localized inside genes; mutation: genes containing predicted editable C \rightarrow T/G \rightarrow A conversion sites; mutation stop: C \rightarrow T/G \rightarrow A conversions generate premature stop codons in genes. (B) Cumulative curve of the distribution of the generated premature stop codons inside gene bodies. (C) Circos plot showing the distribution of genes that could be edited by the pnCasSA-BEC system in the genome of MRSA252. Concentric circles from the inside to the outside: (13) CG content; (12) genes with stop mutation sites; (11) genes with C-T mutation sites; (10) genes with PAM sites; (9) CG offsets ((G - C)/(G + C)); (8-5) genes with stop mutation sites in 0-20%, 20-50%, 50-80%, and 80-100% of the gene body; (4) gene annotations; (3-2) distribution of the genes in forward and reverse strands of the genome; (1) the distribution of the genes in the entire genome.



nickase (Cas9D10A) gene. The D10A mutation in the Cas9 protein was achieved by Gibson assembly.³³ The pCasSA plasmid was separated into two parts. Each part was PCR amplified using the primers containing the desired mutation. The two parts were assembled by Gibson assembly, resulting in the pnCasSA plasmid. The successful construction of the pnCasSA was verified by sequencing.

To construct the pnCasSA-BEC plasmid, we separated the plasmid into four parts. The first fragment contained the Cas9 nickase gene. The second fragment contained the repF origin, the chloramphenicol and kanamycin resistance markers, and the ColE1 origin. The third fragment contained the sgRNA expression cassette. The aforementioned three fragments were PCR amplified from the pnCasSA plasmid, respectively, using the primers listed in Table S4.† The fourth fragment of the *E. coli* codon optimized rat APOBEC1 gene was synthesized by GENEWIZ (Suzhou, China) and PCR amplified using the primers listed in Table S4.† The first and the second fragments were assembled first by Gibson assembly, resulting in the pnCas plasmid. The pnCas plasmid was linearized by *BsaI*. The fourth fragment containing the APOBEC1 gene was inserted into the pnCas plasmid by Gibson assembly, resulting in the pnCas-BEC plasmid. The pnCas-BEC plasmid was linearized by *BsaI* again. The third fragment containing the sgRNA expression cassette was inserted into the pnCas-BEC plasmid by Gibson assembly, resulting in the final plasmid pnCasSA-BEC. The successful construction of the pnCasSA-BEC plasmid was confirmed by PCR, enzyme digestion, and sequencing.

Electroporation

Electrocompetent cells of the *S. aureus* strains were prepared using the following protocol. A single colony of each *S. aureus* strain was inoculated into 2 mL TSB and incubated at 30 °C for 12 h. The cells were 1 : 100 diluted into 100 mL fresh TSB medium. The cells were grown to OD₆₀₀ of ~0.3–0.4 and chilled on ice for 20 min. The cells were harvested by centrifugation (5000 rpm) at 4 °C for 6 min, and washed twice with 20 mL sterile ice-cold 0.5 M sucrose. Finally, the cells were resuspended into 1 mL 0.5 M sucrose. 50 μL aliquots were frozen in liquid nitrogen and stored at –80 °C.

Before electroporation, 50 μL competent cells and the pnCasSA-BEC plasmid were thawed on ice for 5 min. The cells were mixed with 1 μg of the pnCasSA-BEC plasmid, and transferred into a 1 mm electroporation cuvette (Bio-Rad) at room temperature. After being pulsed at 2.1 kV mm⁻¹, 100 Ω, and 25 μF, the cells were immediately supplied with 950 μL TSB and recovered at 30 °C for 1–2 h. The cells were plated onto a TSB plate containing chloramphenicol and incubated at 30 °C for 24 h.

Plasmid curing

The cells containing the desired mutation were cultured in TSB containing 10 μg mL⁻¹ chloramphenicol. They were incubated at 30 °C overnight. The next day, the cells were 1 : 1000 diluted and incubated at 42 °C for 12 h without adding chloramphenicol. The bacteria were plated onto a TSB plate in the absence of

chloramphenicol. The plate was incubated at 37 °C overnight. The next day, several colonies were randomly picked and resuspended individually into 10 μL ddH₂O. The cells were streaked onto two different TSB plates in the presence and absence of 5 μg mL⁻¹ chloramphenicol. The successful curing of the plasmid was confirmed when the cells could only grow on the plate without chloramphenicol.

Hemolysis assay

Various *S. aureus* strains were grown overnight in TSB at 30 °C. The next day, 2 μL cells were loaded onto a sheep blood plate. The plate was incubated at 30 °C for 24 h and further stored at 4 °C for 24 h before being photographed.

Editing efficiency evaluation

The genomic DNA of each colony was purified individually. The PCR reaction was performed individually using the genomic DNA as the template. The PCR products covering the editable site(s) were sent out for sequencing. The editing efficiency was calculated using the following method: the number of PCR products that contained the desired C → T conversion divided by the number of all the PCR products sent out for sequencing.

Bioinformatics analysis

The genome sequences of *S. aureus* MRSA252 and Newman strains were obtained from NCBI (accession numbers: 299279 and 299277). The locations of all PAM (NGG) sites were scanned across the entire genome for the first time. Editable sites with the presence of C(s) at the 13–17 bp upstream of NGG were selected for further analysis. The sites that could introduce stop codons (UAG, UAA, and UGA) after the C → T conversion were treated as editable stop sites. Integrative Genomics Viewer (IGV)³⁴ was used for the visualization of the genes with editable stop sites. Circos (<http://circos.ca>)³⁵ was used for visualizing editable genes of the pnCasSA-BEC system in the genome of the *S. aureus* MRSA252 strain.

Conflicts of interest

A patent has been submitted for the *S. aureus* base-editing method.

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