CRISPR Interference Limits Horizontal Gene Transfer in Staphylococci by Targeting DNA

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Horizontal gene transfer (HGT) in bacteria and archaea occurs through phage transduction, transformation, or conjugation, and the latter is particularly important for the spread of antibiotic resistance. Clustered, regularly interspaced, short palindromic repeat (CRISPR) loci confer sequence-directed immunity against phages. A clinical isolate of *Staphylococcus epidermidis* harbors a CRISPR spacer that matches the nickase gene present in nearly all staphylococcal conjugative plasmids. Here we show that CRISPR interference prevents conjugation and plasmid transformation in *S. epidermidis*. Insertion of a self-splicing intron into nickase blocks interference despite the reconstitution of the target sequence in the spliced mRNA, which indicates that the interference machinery targets DNA directly. We conclude that CRISPR loci counteract multiple routes of HGT and can limit the spread of antibiotic resistance in pathogenic bacteria.

Clustered, regularly interspaced, short palindromic repeat (CRISPR) loci are present in ~40% of eubacterial genomes and nearly all archaeal genomes sequenced to date and consist of short (~24 to 48 nucleotides) repeats separated by similarly sized unique spacers (1, 2). They are generally flanked by a set of CRISPR-associated (cas) protein-coding genes (3–5). The CRISPR spacers and repeats are transcribed and processed into small CRISPR RNAs (crRNAs) (4, 6–8) that specify acquired immunity against bacteriophage infection by a mechanism that relies on the strict identity between CRISPR spacers and phage targets (3, 4).

The rise of hospital- and community-acquired methicillin- and vancomycin-resistant *Staphylococcus aureus* (MRSA and VRSA, respectively) is directly linked to the horizontal transfer of antibiotic resistance genes by plasmid conjugation (9, 10). *S. aureus* and *S. epidermidis* strains are the most common causes of nosocomial infections (11–13), and conjugative plasmids can spread from one species to the other. Although the *S. epidermidis* strain American Type Culture Collection (ATCC) 12228 (14) lacks CRISPR sequences, the clinically isolated strain RP62a (15) contains a CRISPR locus (Fig. 1A and fig. S1A) that includes a spacer (spc1) that is homologous to a region of the nickase (nes) gene found in all sequenced staphylococcal conjugative plasmids (fig. S1B), including those from MRSA and VRSA strains (9, 16, 17).

To test whether spc1 prevents plasmid conjugation into *S. epidermidis* RP62a, we disrupted the sequence match by introducing nine silent mutations into the nes target in the conjugative plasmid pG0400 (18), generating pG0(mut) (Fig. 1B) (19). We tested whether both wild-type and mutant pG0400 transferred from *S. aureus* strain RN4220 (20) into either of the two *S. epidermidis* strains (Fig. 1D and fig. S1C). Although the conjugation frequency of both plasmids was similar for the CRISPR-negative ATCC 12228 strain, only pG0(mut) was transferred into the CRISPR-positive RP62a strain and with a frequency similar to that of wild-type pG0400 in the control ATCC 12228 strain. These results indicate that CRISPR interference can prevent plasmid conjugation in a manner that is specified by sequence identity between a spacer and a plasmid target sequence.

To test this conclusion more rigorously, and to determine whether the CRISPR sequences themselves are responsible for the observed interference, we deleted the four repeats and spacer downstream of an isopropyl-thiogalactopyranoside (IPTG)-inducible pro-

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Fig. 1. A CRISPR locus provides immunity against plasmid conjugation in *S. epidermidis*. (A) Organization of the RP62a CRISPR locus. Repeats and spacers (colored boxes) are followed by CRISPR-associated genes (cas1, cas2, and cas6) and cas subtype Mycobacterium tuberculosis genes (csm1 to csm6) (5). An AT-rich “leader” sequence precedes the repeat-spacer region (black box). LAM104 is an isogenic Δcrispr strain lacking only the repeat and spacer sequences. (B) The staphylococcal conjugative plasmid pG0400 spc1 target sequence [pG0(wt), highlighted in yellow] is shown on the top. This sequence was altered to introduce synonymous mutations to generate pG0(mut), with changes shown in red. (C) To restore interference in strain LAM104, two plasmids were introduced: pCRISPR and pCRISPR-L. (D) Conjugation was carried out by filter-mating in triplicate; the colony-forming units (CFU) per milliliter values (mean ± SD) obtained for recipients and transconjugants are shown. Recipient strains, complementing plasmids, and donor conjugative plasmids are indicated. Conjugation efficiency (Conj. Eff.) was calculated as the recipients/transconjugants ratio.
interference in the spacer/motif, but they differ in the amount of upstream flanking sequence included. The appearance of the spc1 crRNA in pCRISPR-L–containing LAM104 cells (fig. S2) was IPTG-independent, which indicates that the insert includes a natural CRISPR spacer/motif. pCRISPR-L restored interference in the Δcrispr strain, but pCRISPR-L did not, even in the presence of IPTG (fig. S1D). This suggests a role for the leader sequence (Fig. 1A) or other upstream sequences in cis during CRISPR interference, perhaps for processing of the crRNA precursor by Cascade proteins (4). Introduction of pCRISPR-L into strain ATCC 12228 did not alter the conjugation efficiency of pG0400 (fig. S1D) as expected because of the lack of cas genes in this strain.

The nature of the crRNA targeting event (RNA-RNA or RNA-DNA) is not known. The requirement for nickase activity only in the donor cell (16) implies that interference with nes mRNA and protein expression should block the ability of S. epidermidis RP62a to function as a donor. Instead, our data (Fig. 1 and fig. S1) indicate that the CRISPR locus limits the ability of S. epidermidis RP62a to act as a plasmid recipient and therefore suggest that spc1-directed interference does not target the nes transcript. Consistent with this, the orientation of spc1 leads to the expression of a crRNA that is identical with, rather than complementary to, the nes mRNA target sequence, and we have found no evidence for the expression of RNA from the opposite strand (fig. S2) (4). Alternatively, the spc1 crRNA may target the incoming DNA. To test this, we interrupted the target sequence of the pG0400 nes gene with the orf142-II self-splicing group I intron from the staphylococcal Twort phage (21, 22). The mutant conjugative plasmid pG0(I2) lacks an intact spc1 target DNA sequence (Fig. 2A), but the spc1 target sequence is regenerated in the nes mRNA after transcription and rapid splicing. When tested in conjugation assays, pG0(I2) was transferred to wild-type and Δcrispr strains with equal efficiencies (Fig. 2B). This observation reflects an evasion of CRISPR activity when the intron is present in the plasmid and therefore indicates that an intact target site is required in the nes DNA, but not mRNA, for interference to occur. To confirm that splicing is required for nes function, we tested the conjugation ability of pG0(dII), a derivative of pG0(I2) containing a three-nucleotide deletion within the intron that inactivates self-splicing (21) (fig. S3A). The pG0(dII) plasmid was unable to transfer to the RP62a strain (fig. S3B), which indicates a lack of nickase activity in the presence of the unspliced intron.

The requirement for nes transcription, splicing, and translation in the donor cell during conjugation (16), and our ability to obtain RP62a transconjugants with the intron-containing plasmid, allowed us to test the capacity of the CRISPR system to target intact spliced nes mRNA by using RP62a as a pG0(I2) donor. pG0(I2) conjugative transfer was just as efficient from RP62a as from the isogenic Δcrispr strain LAM104 (Fig. 2), which indicates that spliced functional nes mRNA (which must be present for conjugation to occur) is not targeted during CRISPR interference. Reverse transcription polymerase chain reaction assays confirmed the splicing of the nes premRNA in RP62a cells carrying pG0(I2) (fig. S3C). Although our observations do not formally exclude an RNA targeting event that is somehow restricted to nascent, transient, unspliced transcripts, they provide strong evidence that DNA rather than mRNA is the likely crRNA target during CRISPR interference.
CRISPR activity against phage and conjugative plasmid DNA molecules suggests that CRISPR systems may also prevent plasmid DNA transformation. We therefore introduced pG0(wt) and pG0(mut) nes-target and -flanking sequences (200 base pairs) in either orientation into the staphylococcal plasmid pC194 (23), generating pNes(wt) and pNes(mut), respectively (Fig. 3A). Flanking DNA was included in the inserts to ensure the presence of any sequences outside of the target that may contribute to CRISPR interference (24). Plasmids were transformed by electroporation into wide-type RP62a and isogenic Δcrispr LAM104 strains. pC194 and both pNes(mut) plasmids were transformed into both strains, whereas the pNes(wt) plasmids were transformed only into the Δcrispr mutant (Fig. 3B). We also performed pNes(wt)/pNes(mut) mixed transformations of RP62a or LAM104 strains to test interference in an internally controlled fashion. Again, only pNes(mut) plasmids were recovered from RP62a transformants, whereas pNes(wt) and pNes(mut) plasmids were found in LAM104 transformant colonies (fig. S4). It remains to be established whether natural transformation, which involves the uptake of a single DNA strand (25), is subject to CRISPR interference. Nonetheless, our experiments suggest that CRISPR systems can counteract multiple routes of plasmid transfer.

These transformation data provide additional evidence that crRNAs target DNA molecules. First, interference occurred regardless of the insert orientation in pNes(wt); this, combined with the lack of compelling evidence for CRISPR-derived double-stranded RNA (fig. S2) (4, 6, 7), is consistent with spec1 targeting either DNA strand rather than a unidirectional transcript. Second, the target sites in the pNes(wt) and pNes(mut) plasmids are located between the transcriptional terminators of the rep and cat genes (Fig. 3A) (23, 26, 27). This minimizes the likelihood that this region of the plasmid is even transcribed, which is consistent with its dispensability for plasmid maintenance (23, 28).

Altogether, these data provide strong functional evidence that CRISPR interference acts at the DNA level and therefore differs fundamentally from the RNA interference (RNAi) phenomenon observed in eukaryotes and with which CRISPR activity was originally compared (29). A DNA targeting mechanism for CRISPR interference implies a means to prevent its action at the encoding CRISPR locus itself, as well as other potential chromosomal loci, such as prophage sequences. Little information exists to suggest how crRNAs would avoid targeting “self” DNA, although the role of flanking sequences during CRISPR interference (24) could contribute to target specificity. From a practical standpoint, the ability to direct the specific addressable destruction of DNA that contains any given 24- to 48-nucleotide target sequence could have considerable functional utility, especially if the system can function outside of its native bacterial or archaenal context. Furthermore, our results demonstrate that CRISPR function is not limited to phage defense, but instead encompasses a more general role in the prevention of HGT and the maintenance of genetic identity, as with restriction-modification systems. A primary difference between restriction-modification and CRISPR interference is that the latter can be programmed by a suitable effector crRNA. If CRISPR interference could be manipulated in a clinical setting, it would provide a means to impede the ever-worsening spread of antibiotic resistance genes and virulence factors in staphylococci and other bacterial pathogens.

References and Notes
19. Materials and methods are available as supporting material on Science Online.
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Supporting Online Material
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Materials and Methods
Figs. 51 to 54
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References
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Nascent RNA Sequencing Reveals Widespread Pausing and Divergent Initiation at Human Promoters
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RNA polymerases are highly regulated molecular machines. We present a method (global run-on sequencing, GRO-seq) that maps the position, amount, and orientation of transcriptionally engaged RNA polymerases genome-wide. In this method, nuclear run-on RNA molecules are subjected to large-scale parallel sequencing and mapped to the genome. We show that peaks of promoter-proximal pausing reside on ~30% of human genes, transcription extends beyond pre-messenger RNA 3′ cleavage, and antisense transcription is prevalent. Additionally, most promoters have an engaged polymerase upstream and in an orientation opposite to the annotated gene. This divergent polymerase is associated with active genes but does not elongate effectively beyond the promoter. These results imply that the interplay between polymerases and regulators over broad promoter regions dictates the orientation and efficiency of productive transcription.

Transcription of coding and noncoding RNA molecules by eukaryotic RNA polymerases requires their collaboration with hundreds of transcription factors to direct and control polymerase recruitment, initiation, elongation, and termination. Whole-genome microarrays and ultra-high-throughput sequencing technologies enable efficient mapping of the distribution of transcription factors, nucleosomes, and their modifications, as well as accumulated RNA transcripts throughout genomes (1, 2), thereby providing a global correlation of factors and transcription states. Studies using the chromatin immunoprecipitation assay coupled to genomic DNA microarrays (ChIP-chip) or to high-throughput sequencing (ChIP-seq) indicate that RNA polymerase II (Pol II) is present at disproportionately higher amounts near the 5′ end of many eukaryotic
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