
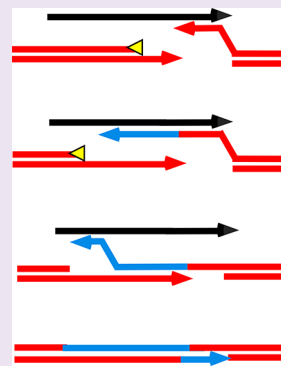


# Repair of a Site-Specific DNA Cleavage: Old-School Lessons for Cas9-Mediated Gene Editing

Danielle N. Gallagher and James E. Haber\*

Rosenstiel Basic Medical Sciences Research Center and Department of Biology, Brandeis University, Waltham, Massachusetts 02454-9110, United States

**ABSTRACT:** CRISPR/Cas9-mediated gene editing may involve nonhomologous end-joining to create various insertion/deletions (indels) or may employ homologous recombination to modify precisely the target DNA sequence. Our understanding of these processes has been guided by earlier studies using other site-specific endonucleases, both in model organisms such as budding yeast and in mammalian cells. We briefly review what has been gleaned from such studies using the HO and I-SceI endonucleases and how these findings guide current gene editing strategies.



## ■ HO AND I-SCEI ENDONUCLEASES

The HO and I-SceI are members of a large family of evolutionarily related site-specific endonucleases that create double-strand breaks (DSBs) with 4-nt 3' overhangs.<sup>1,2</sup> In budding yeast, these enzymes provoke specific chromosome rearrangements in which sequences from a donor locus are copied by a process known as gene conversion to patch up the broken chromosome.

The HO endonuclease causes the switching of yeast mating-types, from *MATa* to *MATα* or vice versa<sup>3,4</sup> (Figure 1A). *MAT* switching involves the removal of about 700 bp of mating type-specific sequences (Ya or Yα) and their replacement by the opposite sequences that are copied from one of two heterochromatic and silent donor loci, *HMLα* and *HMRα*, that are each located >100 kb away from *MAT* on the same chromosome. HO endonuclease (hereafter referred to as HO) efficiently cleaves a degenerate 24-bp sequence, so that either *MATa* and *MATα* sequences are cut; the same recognition sites in *HMLα* and *HMRα* are normally not cut because these heterochromatic loci have highly positioned nucleosomes that apparently block cleavage. Cleavage at the donors is efficient only when the silencing, via the Sir2 histone deacetylase, is disabled. The HO cleavage site can be inserted into any other euchromatic locus and is often employed to study various mechanisms of DNA repair (discussed below). At its normal location within the *MAT* locus or when inserted elsewhere, the HO cut site is readily cleaved when HO endonuclease is induced, by placing the HO gene under the control of a galactose-inducible promoter.<sup>5</sup> Cleavage is quite rapid, with >90% of the cells being cleaved in 30–45 min, leaving 4-nt 3'-overhanging DSB ends.<sup>6,7</sup> This makes it possible to monitor in real time the sequence of events leading to the formation of a repaired DSB for several different types of DSB repair by

homologous recombination<sup>8–16</sup> (Figure 2), as well as by nonhomologous end-joining (NHEJ).<sup>17</sup>

Very little use of HO has been made in metazoans, as the enzyme has never been characterized in detail *in vitro*, unlike I-SceI.<sup>18–20</sup>

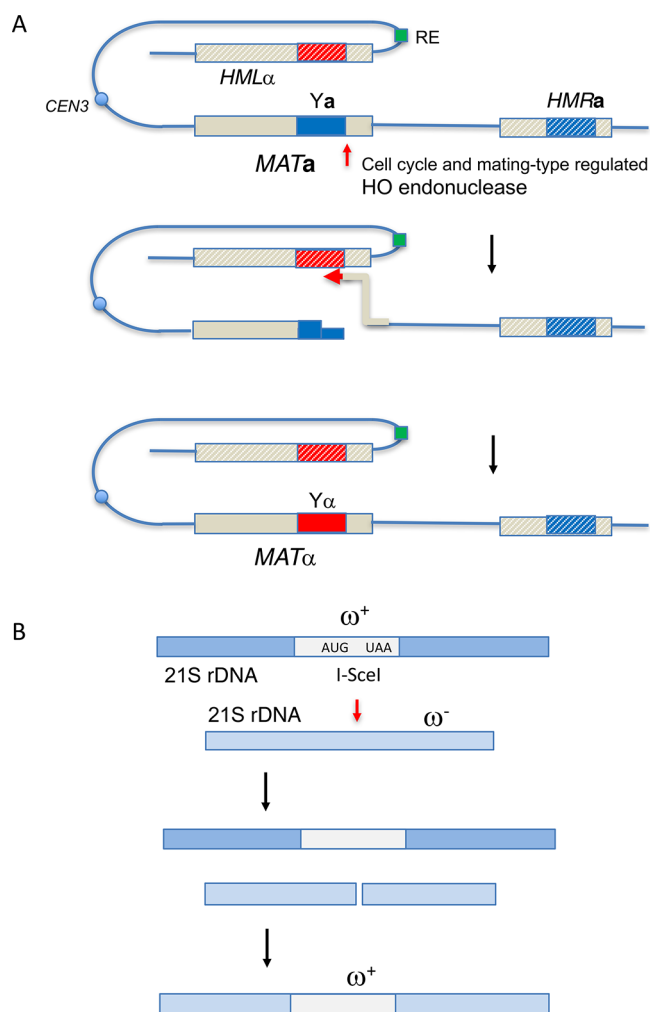
**I-SceI in Yeast and Mammals.** I-SceI is normally localized in the budding yeast mitochondrion, encoded within an intron within the 21S rDNA. This intron is known as omega and designated  $\omega^+$ . When an  $\omega^+$  strain is crossed to an  $\omega^-$  strain, there is a very efficient conversion of the  $\omega^-$  mitochondrial genome to the intron-containing  $\omega^{+21,22}$  by creating a site-specific DSB within the  $\omega^-$  21S rDNA sequence and then using the  $\omega^+$  copy as the repair template, through homologous recombination. I-SceI cannot cleave the inserted  $\omega^+$  sequence. This example of “intron homing” was the first well-studied example of gene drive, as all the many copies of mitochondrial DNA in progeny all become  $\omega^+$  (Figure 1B). I-SceI was “domesticated” for general use—to study chromosomal repair in the nucleus—by recoding the gene and expressing it in the nucleus so that it was compatible with cytoplasmic protein synthesis.<sup>23</sup> I-SceI has an 18-bp recognition site. Placed under the same galactose-inducible promoter as in the case of HO, I-SceI cleavages are reasonably efficient (though less than HO), and repair can be monitored both physically and genetically.<sup>24</sup> I-SceI also makes 4-nt, 3'-overhanging ends and both the homologous recombination events and NHEJ outcomes are comparable to what had been shown for HO.<sup>18</sup> The I-SceI cleavage site can be inserted in different locations to study

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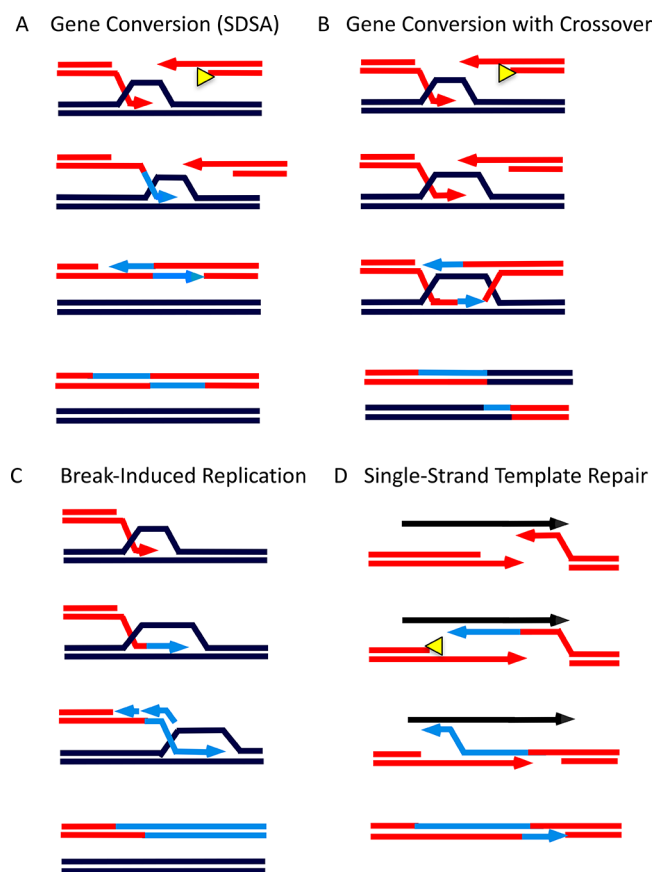
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**Figure 1.** HO and I-SceI-mediated recombination in budding yeast. (A) Mating-type (*MAT*) gene switching. Prior to S phase, budding yeast express HO endonuclease, resulting in the cleavage of either *MAT $\alpha$*  or *MAT $\beta$*  on chromosome 3. The silent, heterochromatic structure (hatched lines) of the two donors, *HML $\alpha$*  and *HMR $\alpha$* , located 200 and 100 kb from *MAT*, respectively, are resistant to cleavage but serve as efficient donors to repair the DSB. Donor preference is regulated by the cis-acting Recombination Enhancer (RE); in *MAT $\alpha$*  cells, proteins bound to RE also bind near the DSB end and promote the use of *HML*. In *MAT $\beta$* , RE is itself silenced, allowing *HMR $\alpha$*  to be the default donor. Repair of the DSB involves removal of the original mating type-specific *Y $\alpha$*  or *Y $\beta$*  sequences and their replacement with the opposite mating type. DSB repair by gene conversion without an accompanying exchange (see Figure 2A) is the predominant mechanism of repair. (B) Intron homing of the omega  $\omega^+$  intron within the mitochondrial 21S rDNA. When an  $\omega^+$  budding yeast mates with an  $\omega^-$  cell of opposite mating type, I-SceI endonuclease, encoded within the intron, cleaves its target site in the  $\omega^-$  locus and promotes repair by gene conversion. Details of the molecular events that lead to repair have not been established in detail. As there are often multiple copies of each mitochondrial genome, the global replacement of  $\omega^-$  by  $\omega^+$  is the first example of gene drive.

various aspects of homologous recombination in budding yeast.<sup>25–30</sup>

In mammalian cells, I-SceI became the principal means to induce site-specific DSBs and to learn about DSB repair in detail.<sup>31,32</sup> Although site-specific nucleases (zinc-finger nucleases, TALENs, and of course CRISPR/Cas9) have emerged to facilitate gene targeting, most of the mechanistic under-



**Figure 2.** Mechanisms to repair a double-strand break. (A) Gene conversion by synthesis-dependent strand annealing (SDSA) allows repair by patching up a DSB without chromosomal rearrangement. Key steps include 5' to 3' resection of DSB ends, recruitment of Rad51 recombinase as a filament on the 3'-ended single-stranded DNA (ssDNA), a homology search culminating in the formation of a displacement loop (D-loop) by base-pairing between the invading strand and the complementary homologous strand in the donor locus, initiation of new DNA synthesis, displacement of the newly synthesized DNA to anneal with the second end of the DSB (second end-capture), and completion of repair by second-end DNA synthesis and ligation of the strands. (B) Gene conversion via a double-Holliday junction (dHJ) intermediate follows the same initial steps, but an extended D-loop is annealed by the second end. New DNA synthesis and ligation lead to a dHJ structure that can be “dissolved” to a noncrossover outcome or resolved by structure-specific nucleases to produce a crossover outcome. (C) Break-induced replication (BIR) begins similarly to SDSA, but new DNA synthesis requires all of the normal replication apparatus plus at least two proteins not needed for gene conversion or normal DNA replication. Although both leading- and lagging-strand synthesis must occur, they are not coordinated as in normal replication, and all the newly copied DNA is conservatively inherited at the initially broken end. (D) In single-strand template repair (SSTR), only one resected end of the DSB can pair with the ssDNA template and initiate new DNA synthesis, but there is apparently no need for Rad51 and strand invasion to accomplish this step. The newly extended DNA is apparently displaced from the template and can anneal with the second end—analogue to second-end-capture—and second end synthesis can complete the repair of the DSB.

standing of DSB repair in mammals has continued to rely on I-SceI. Maria Jasin's lab led the way in developing assays for both homologous recombination and NHEJ, demonstrating that both gene conversions and NHEJ-mediated indels could be

recovered after I-SceI cleavage.<sup>33–37</sup> Many laboratories have contributed to a number of different assays that make it possible to study gene conversions, SSA, and end-joinings.<sup>38–42</sup> Assays that mimic break-induced replication (BIR) have also been developed.<sup>43</sup> One important distinction between yeast and mammals is that interchromosomal ectopic recombination events are quite rare. However, recovery of reciprocal translocations from two DSBs on different chromosomes—one pair of ends using single-strand annealing (SSA) and the other pair using NHEJ—can readily be recovered.<sup>44–46</sup> Also, it should be noted that, compared to budding yeast, NHEJ is much more efficient relative to homologs in mammalian cells; however, often, about half of all detectable recombination/repair events in Cas9 or I-SceI-mediated repair are still *via* homologous recombination.<sup>47–49</sup> Another distinction between events in yeast and those in mammals is that, even when both sides of the DSB share homology with a donor sequence, many events begin as homologous recombination but terminate by some sort of NHEJ event. *The capacity of mammalian cells to shift from one mode of repair to the other may pose problems in the fidelity of gene editing.*

## ■ DEFINING AND MONITORING MULTIPLE PATHWAYS OF DNA REPAIR IN YEAST

**DSB Repair by Gene Conversion.** *MAT* switching is perhaps the best-studied example of double-strand break repair that is rarely associated with crossing-over; one set of sequences at *MAT* is replaced by another, copied from a donor. The synchronous cleavage by HO allowed us to identify a number of surprisingly slow steps in DSB repair (Figure 2A), including:

(a) 5' to 3' resection of DNA ends by several exonucleases, including the Mre11–Rad50 complex, ExoI, and Sgs1–Rmi1–Top3–Dna2, as well as the chromatin remodeler, Fun30, which interacts with nucleosomes proximal to the DSB to facilitate exonuclease digestion of the DNA (illustrated by the yellow triangle in Figure 2A)<sup>14,50–56</sup>

(b) assembly of single-strand DNA binding protein RPA on the 3'-ended single-strand DNA (ssDNA), facilitating the formation of the Rad51 filament that will search the genome for a homologous region<sup>57</sup>

(c) rapid association of Rad51 with the donor sequence and the slower formation of base pairs between the invading 3' ssDNA with the dsDNA donor, creating a displacement loop (D-loop; Figure 2A, top line)<sup>58,59</sup>

(d) initiation of new DNA synthesis, principally by DNA polymerase  $\delta$  (Figure 2A, second line, blue arrow)<sup>14,60</sup>

(e) capture of the newly copied and displaced DNA by the homologous sequences of the second DSB end (Figure 2A, third line), by strand annealing of the newly copied DNA with the resected second end<sup>61,62</sup>

(f) completion of repair, which takes about 1–2 h; the sequence of DNA synthesis events results in all the newly copied DNA in the recipient locus, and the donor is unchanged (Figure 2A, bottom line)<sup>63</sup>

More detailed descriptions of the molecular mechanisms of this and other repair processes can be found in recent reviews.<sup>4,16</sup>

Interchromosomal DSB repair is also accompanied by crossing-over in a small fraction of cases, as detected by novel restriction fragments on Southern blots.<sup>64</sup> The proportion of crossovers is markedly increased by deleting components of a double-Holliday junction “dissolvase” (the 3' to 5' helicase Sgs1, homologous to human BLM or

Topoisomerase 3 $\alpha$ ).<sup>65,64</sup> Thus, intermediates are frequently formed that could lead to reciprocal exchanges (Figure 2B), but these are usually defeated by their dissolution. Crossovers also increase after deleting another 3' to 5' helicase, Mph1 (related to human FANCM).<sup>66</sup> Mph1 apparently directs repair to the noncrossover synthesis-dependent strand annealing (SDSA) pathway described above and away from double-Holliday junction intermediates.

A surprising discovery was that—although gene conversion events are the most faithful way to repair a DSB, without the alterations that would arise by crossing-over and also without the creation of insertion/deletions by nonhomologous end-joining—the rate of mutation in the newly copied sequences is 1000 times that of normal replication.<sup>67–69</sup> Many of the mutations result from DNA polymerase  $\delta$  errors, including simple DNA polymerase slippage errors such as –1 frameshifts in homonucleotide sequences (i.e., CCCC becoming CCC) but including more dramatic interchromosomal template switches between highly divergent sequences.<sup>70</sup> *Whether mammalian gene editing is likewise associated with such errors remains to be learned.*

**Break-Induced Replication (BIR).** When only one end of a DSB is homologous to a donor sequence, repair shifts to a recombination-dependent initiation of DNA synthesis that—at least in yeast—can copy several hundred kilo-base pairs to the end of a chromosome, resulting in a nonreciprocal translocation<sup>8,71,72</sup> (Figure 2C). Such events are believed to occur naturally in the restart of stalled and broken DNA replication forks or at eroded telomeres. When both ends of a DSB share homology with a donor sequence, gene conversion overshadows BIR by the agency of a recombination execution checkpoint that delays BIR and apparently allows cells the opportunity to complete gene conversion instead.<sup>73</sup>

Although BIR appears to be the establishment of a normal leading/lagging-strand replication fork, evidence from yeast argues that the synthesis of the two strands is dis-coordinated, with a long single-strand tail being generated behind a moving replication bubble copying the leading strand; only later is the lagging strand added.<sup>74</sup> As in SDSA, all the newly copied DNA is found in the recipient, and there is again a 1000-fold increase in mutations, with a great sensitivity to agents that deaminate or modify the ssDNA before it is made double-stranded.<sup>75</sup> The replication process in BIR requires two proteins that are not essential for normal replication, the Pol $\delta$ -associated Pol32 protein and the 5' to 3' helicase Pif1.<sup>76–78</sup> A dominant mutation in PCNA also blocks BIR without affecting gene conversion or S-phase replication.<sup>79</sup>

In fact, there are two BIR processes. There is an efficient Rad51-dependent event that requires the Rad55/57 paralogs and the Rad54 chromatin remodeler, but there is also a much less efficient Rad51-independent BIR process.<sup>78</sup> Rad51-independent BIR requires the strand annealing protein Rad52, as well as its “cousin,” Rad59, which promotes strand annealing. This pathway also requires the Mre11–Rad50–Xrs2 (MRX) complex and Rdh54 (a homolog of Rad54). Rad51-dependent repair is strongly Rad54-dependent but Rdh54-independent, although error-prone template switching events during gene conversion do rely on Rdh54. How these proteins would cooperate to initiate new DNA synthesis to repair the DSB end is not at all evident. This Rad51-independent pathway also acts preferentially when only short homologous sequences are available, including at yeast's highly irregular telomeres.<sup>78,80</sup> The nonessential Pol32 protein is also needed for Rad51-



independent telomere maintenance without telomerase.<sup>76</sup> An important question to be considered is whether Rad51-independent single-strand template repair (SSTR) in gene editing resembles this Rad51-independent process or represents an entirely different mechanism.

**Single-Strand Annealing (SSA).** The simplest homologous recombination process is single-strand annealing (SSA) where homologous sequences flanking a single DSB are resected to reveal complementary single strands that can anneal, after which nonhomologous 3'-ended tails need to be clipped off and the remaining single-stranded regions are filled in. In budding yeast, 1 kb sequences that are  $\geq 25$  kb can be annealed in this way, creating large deletions.<sup>73,81</sup> Interchromosomal rearrangements can be achieved by creating DSBs on two different chromosomes, with appropriate adjacent homologous sequences to force reciprocal translocations.<sup>82</sup> These processes are Rad51-independent but require the Rad52 and Rad59 strand annealing proteins, as well as an Slx4, Rad1–Rad10, and Msh2–Msh3-dependent apparatus to clip off the nonhomologous tails.<sup>83–88</sup> How terminal nonhomologies are dealt with in SSTR is also not known.

**Nonhomologous End-Joining (NHEJ) and Microhomology-Mediated End-Joining (MMEJ).** "Classic" NHEJ (c-NHEJ) is highly conserved from yeast to humans, requiring the Ku70 and Ku80 proteins, DNA ligase 4, and its associated XRCC4.<sup>89–91</sup> Another Dnl4-associated protein, Nej1, is surprisingly divergent among species, sharing only a few limited motifs with its presumed mammalian homolog, XLF/Cernunnos.<sup>92</sup> Most NHEJ events in budding yeast require the MRX complex,<sup>17</sup> whereas the mammalian Mre11–Rad50–Nbs1 (MRN) complex appears to be most important for alternative end-joining.<sup>93,94</sup>

Alternative end-joining (also called microhomology-mediated mediated end-joining (MMEJ)), is a hybrid pathway which also entails annealing between very short sequences exposed by exonucleases, but MMEJ is distinct from SSA and from c-NHEJ.<sup>95,96</sup> MMEJ is independent of Rad52; whether Rad59 is required needs to be tested. Unlike SSA, MMEJ in budding yeast is partially dependent on DNA ligase4, but unlike c-NHEJ, this process is Ku-independent. In mammals, alternative end-joining is independent of Ku, DNA ligase IV, and XRCC4 but is impaired both by inhibiting poly(ADP-ribose) polymerase (PARP) and DNA ligase III (both of which are absent in budding yeast). Because many "classic" (Ku-dependent) NHEJ events are found to have a few bases of microhomology at the junctions of deletions,<sup>97</sup> it is difficult to specify precisely how much microhomology is required to result in MMEJ, but experimental models have used microhomologies of 6 or more bases.<sup>98</sup>

When HO is induced and then shut off, the enzyme is rapidly degraded, and one can visualize rejoining of the 4-nt, 3' overhanging ends to reform the original cleavage site.<sup>99</sup> In a strain lacking homologous donors, simple religation of the overhanging ends is an efficient process, requiring Ku proteins, Dnl4, and other parts of the c-NHEJ machinery.<sup>17</sup> Whether Cas9-mediated perfect rejoins have the same genetic requirements is not yet known, because it is necessary both to induce synchronous cleavage and then to turn off further enzyme activity. Alternatively, one could assay the blunt-end joining of two efficiently cleaved sites.<sup>100</sup> Such assays have not been done yet to look at the kinetics of the process.

If HO expression is maintained, resection of the ends renders precise end-joining less and less efficient. In the face of

continuous HO expression, survivors emerge; these have mutated the cleavage site and therefore can no longer be cut. Insertions and deletions (indels) predominate over single base-pair mutations. The great majority of these indels result from various mispairings of the 4-nt overhanging ends to produce 2 and 3 bp insertions and 3 bp deletions.<sup>17</sup> These events all involve single base pairings, trimming overhanging unpaired ends, and the filling in the small gaps, by the PolX polymerase, Pol4. Other deletions of varying size are recovered at lower rates. Importantly, the overall spectrum of NHEJ events created by HO is not different from that created by inducing chromosome breakage using a conditionally dicentric chromosome.<sup>101</sup>

**The Age of CRISPR/Cas9.** Although CRISPR/Cas9 has been studied biochemically and *in vivo*, there are still a number of questions that we wish to answer. Does Cas9 stay bound persistently to one end of the DSB, blocking its resection? Given that at least most Cas9 cleavages leave blunt ends, how efficient are their rejoins? How long-lived is the protein::gRNA complex? What is the mechanism of gene editing by single-strand template repair (SSTR)? To monitor such events *in vivo*, it is first necessary to have synchronous cleavage of the target sequence. In mammalian systems, the highest level of cleavage has been accomplished by introducing an *in vitro*-assembled ribonucleoprotein particle (RNP), but this approach has not yet lent itself to following a detailed kinetics of repair, as electroporation is followed by an uncharacterized period of recovery. In budding yeast, it has been possible to accomplish nearly complete and rapid (within 1–2 h) cleavage using an inducible gRNA and constitutively expressed protein, as developed by Smith and co-workers.<sup>102</sup> Several strategies have been used to promote Cas9 activity in a synchronous fashion.<sup>103,104</sup>

How persistent is enzyme activity after induction is turned off? To address this question, one needs to induce cleavage of a conditionally accessible site. Neither HO nor a Cas9 directed to the same location will cleave a sequence inside the heterochromatic *HML $\alpha$*  locus, but cleavage is efficient once silencing is blocked, by inhibiting the Sir2 histone deacetylase with nicotinamide. Hence, it should be possible to ask how long the enzyme is still active, after induction is stopped, by examining cleavage after the desilencing of the target at 2, 4, and 6 h. In mammalian cells, Cas9 cleavage is also dependent on the chromatin context of the target sequence.<sup>105,106</sup> Cleavage is apparently impaired in heterochromatic regions (although a small fraction of sites in major satellite DNA are apparently cleaved).<sup>107</sup> It might be possible to enhance cleavage in such regions by using histone deacetylase inhibitors or other means to block heterochromatin formation.

Because Cas9 is long-lived, ends that are joined can be recut, making it difficult to assess the efficiency of end-joining of the cut locus; hence it is necessary to inactivate the Cas9 protein. This can be accomplished through the use of an auxin-inducible degron,<sup>108</sup> as has also been done for the site-specific AsiSI endonuclease in mammalian cells.<sup>109,110</sup> Another way to destabilize Cas9 has been to fuse it to the FKBP12-L106P domain.<sup>100</sup> In this way, one can observe in budding yeast that the cleaved ends rejoin in real time, as was first done with HO. Whether this can be done in mammals remains to be seen. It should also be noted that there is substantial evidence in mammals<sup>111</sup> and in yeast<sup>112</sup> that some fraction of Cas9 cleavages *in vivo*, as well as *in vitro*,<sup>113</sup> are not blunt-ended but leave a 1-nt 5' overhang that can be filled in (in yeast by a PolX polymerase, Pol4) and then the

new blunt ends joined to create a templated +1 base pair insertion.<sup>112</sup> Whether the overhanging ends are more rapidly rejoined *in vivo* will be challenging to determine.

Efficient cleavage also allows us to assess the resection of the DSB ends by exonucleases. *In vitro* studies have suggested that Cas9 remains asymmetrically bound to one end.<sup>114</sup> Using the same strategies we had developed for measuring resection of HO-cut ends, it should be possible to assess whether one end is retarded in its resection. Whether removal of Cas9 from an end will require the activity of the Mre11 complex (MRX) as do covalently attached ends<sup>115</sup> can readily be determined.

**SSTR versus Other Mechanisms of DSB Repair.** Single strand template repair (SSTR) appears to be a novel process, distinct from other homologous repair events (Figure 2D). It is possible that this pathway was intended to repair DNA *via* single-stranded RNA templates<sup>116</sup> but has now been exploited to facilitate use of ssDNA repair templates provided in excess, along with the programmed endonuclease.<sup>48,117–122</sup> Whether induced by the HO endonuclease or CRISPR/Cas9, a DSB can be repaired (and gene-edited) by single stranded DNA oligonucleotides. SSTR is possible even if the region to be modified is encoded on several similarly oriented oligonucleotides, each partially overlapping the homology on the chromosome or an ssDNA template.<sup>122</sup> Budding yeast is particularly adept in this type of gene editing, as it is possible to create, at efficiencies on the order of 80% of all transformants, simple mutations such as base-pair substitutions, deletions of varying size, or insertions of epitope tags, fluorescent protein fusions, or degrons.<sup>123</sup> Models of SSTR share some similarities with both SDSA and BIR mechanisms that engage double-stranded DNA (dsDNA templates), but SSTR has proven to occur independently of the Rad51 strand exchange protein that is required for almost all dsDNA templates.<sup>119</sup> Storici's initial observation in budding yeast was extended to mammals in the context of repairing DNA nicks produced by the CRISPR/Cas9<sup>D10A</sup> nickase<sup>121</sup> and has now been found for SSTR in both worms and mammals.<sup>48,122</sup>

As noted above, only one Rad51-independent DSB repair pathway has been identified before, a rather inefficient Rad51-independent variant of BIR.<sup>11,78</sup> Rad51-independent BIR depends on Rad52 and Rad59, the MRX complex (MRN in mammals), Rdh54 (a Swi2/Snf2 chromatin remodeler), and Srs2, a 3' to 5' helicase that acts as a Rad51 antagonist. Our initial studies have indicated that SSTR has distinctly different requirements from Rad51-independent BIR and hence is a new and still largely uncharacterized pathway. *A resected DSB end can apparently anneal with ssDNA without the agency of Rad51. How the second end is captured, in both SSTR and Rad51-independent BIR, and how homology is recognized without the canonical homology recognition protein Rad51 remains unanswered.*

The incredibly rapid development of CRISPR/Cas9 in gene editing has been made possible by relying on a long and rich history of studying DSB repair using a few very site-specific endonucleases. There are still important lessons to be learned by studying these earlier paradigms. The ability in yeast to create DSBs in a highly synchronous fashion and to follow their repair in real time provides an opportunity to address a number of outstanding issues concerning Cas9 and to provide a guide for determining if these processes are indeed preserved in mammals, as has been the case for many aspects of DSB repair.<sup>4</sup> Given that Cas9 evolved in a noneukaryotic context, it is likely that the rules of DSB repair and gene targeting in eukaryotes

will generally be independent of the source of cleavage, but the special features of Cas9 or other Cas proteins may yield surprises.

## AUTHOR INFORMATION

### Corresponding Author

\*E-mail: [haber@brandeis.edu](mailto:haber@brandeis.edu).

### ORCID

James E. Haber: 0000-0002-1878-0610

### Notes

The authors declare no competing financial interest.

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## KEYWORDS

**Gene editing:** Alteration of a gene by homologous recombination to change a protein open reading frame or adjacent regulatory sequences

**Homologous recombination:** A set of replication-dependent processes to repair a double-strand chromosome break (DSB) that are mediated by copying identical or similar DNA sequences from an intact template that shares homology with one or both sides of the break

**Single-strand template repair (SSTR):** A process of gene editing in which a site-directed double-strand break on a chromosome is repaired by copying sequences from a single-stranded DNA sequence that alters the guide RNA recognition sequence and may introduce other changes as well; gene alterations may include single base alterations, deletions, or insertions of epitopes or other additional protein motifs

**Gene conversion (GC):** A process of homologous recombination in which both ends of a double-strand chromosome break are able to recombine with a donor template, patching up the break by copying the sequences from the donor, which can be located on a sister chromatid, on a homologous chromosome, or at an ectopic location; gene conversion can occur with or without crossing over

**Break-induced replication (BIR):** A process of homologous recombination in which homology near one end of a double-strand break recombines with a template and establishes a modified unidirectional DNA replication fork that can extend hundreds of kilobase pairs, to a chromosome end; repair can occur between homologous chromosomes or between sequences located at ectopic locations, resulting in non-reciprocal translocations

**CRISPR/Cas9:** An RNA-directed, site-specific endonuclease derived from *Streptococcus pyogenes* that has been widely used to create double-strand chromosome breaks in a wide variety of organisms, allowing both the creation of small insertion/deletion mutations as well as template-directed gene editing

**HO endonuclease:** A site-specific endonuclease, recognizing a degenerate 24-base pair sequence, expressed in budding yeast cells to initiate a homologous recombination process that results in mating-type switching

**I-SceI endonuclease:** A site-specific endonuclease expressed in the mitochondria of budding yeast cells to initiate a homologous recombination process that results in the insertion of an intron containing the DNA sequence of the endonuclease; a synthetically created version of the I-SceI gene has been used to induce site-specific double-strand breaks at an 18-base pair recognition site, in both budding yeast and metazoan

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