Alignment of recombination sites in Hin-mediated site-specific DNA recombination

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The Hin site-specific recombination system normally promotes inversion of DNA between two recombination sites in inverted orientation. We show that the rate of deletion of DNA between two directly repeated recombination sites is 10–300 times slower than inversion between sites in their native configuration as measured in vivo and in vitro, respectively. In vitro studies have shown that the deletion reaction has the same requirement for Fis, a recombinational enhancer, and DNA supercoiling as the inversion reaction. These requirements, together with the finding that the deletion products are interlinked once suggest that the deletion synaptic complex is similar to the invertasome intermediate that generates inversion. The inefficiency of the deletion reaction is not a function of a reduced ability to recognize or synapse recombination sites in direct orientation. Not only do these substrates support an efficient knotting reaction, but directly repeated recombination sites with symmetric core sequences also invert efficiently. These findings demonstrate that the recombinational enhancer promotes inversion regardless of their starting orientation. We propose that the dynamics of a supercoiled DNA molecule biases the geometric assembly of specific intermediates. In the case of Hin-mediated recombination, inversion is overwhelmingly preferred over deletion because DNA supercoiling favors a specific alignment of DNA strands in the synaptic complex.

[Key Words: Hin; Fis; recombinational enhancer; site-specific recombination; DNA strand exchange; DNA supercoiling]

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The product of Hin-mediated site-specific recombination is the inversion of a 955-bp DNA segment in the Salmonella chromosome. Inversion of DNA between the two recombination sites switches the orientation of the promoter that transcribes the H2 flagellin gene and the repressor of the unlinked H1 flagellin gene, resulting in the alternate expression of the H1 and H2 flagellins (Silverman and Simon 1980; Zieg and Simon 1980). Scott and Simon (1982) demonstrated that Hin-mediated recombination in vivo resulted in deletion of the intervening DNA when the orientation of one of the recombination sites is reversed on a plasmid substrate. However, deletion of DNA located between the directly repeated recombination sites occurred much less efficiently than inversion between recombination sites in their native orientation. The related Gin and Cin recombinases invert a segment of DNA containing tail fiber genes in phage Mu and P1, respectively (Kamp et al. 1978; Iida et al. 1982). In each of these systems, the inversion reaction is much more efficient than the deletion reaction (Kennedy et al. 1983; Plasterk et al. 1983). This bias in the relative efficiency of the two reactions is one of the hallmarks of the DNA invertase family.

The Hin-mediated DNA inversion reaction occurs efficiently in vitro with purified components (Johnson et al. 1986). Inversion requires a supercoiled DNA substrate containing the two recombination sites (hix sequences) and a recombinational enhancer sequence (Johnson and Simon 1985). The recombinational enhancer is a 63-bp DNA segment that is located 99 bp from the left recombination site in its native context but can function many kilobases away from either recombination site to promote inversion. Each recombination site consists of a 26-bp sequence in which two imperfect 12-bp half-sites are separated by a 2-bp core where strand exchange occurs (Johnson and Simon 1985; Johnson and Bruist 1989). The relative orientation of the recombination sites with respect to each other is defined by their configuration in the Salmonella chromosome and the products of recombination. The hix sites in their native configuration (inverted) give rise to inversion of the intervening DNA, whereas in vivo analysis has shown that when one site is
oriented in the reverse configuration (directly repeated) the intervening DNA segment is deleted (Scott and Simon 1982). Because of the symmetrical nature of the hix sequence, the molecular features that determine directionality of the site have been enigmatic.

Three proteins, Hin, Fis, and HU, are required for efficient inversion in a wild-type plasmid substrate (Johnson et al. 1986). The Hin recombinase binds to each recombination site and catalyzes a staggered cleavage leaving the 2-bp core nucleotides as 3'-single-stranded ends. Hin remains covalently associated with the 5' end of the cleavage site until the DNA is ressealed (Johnson and Bruist 1989). Fis is a 98-amino-acid polypeptide that was first identified by its role in promoting site-specific DNA inversion (Johnson et al. 1986; Koch and Kahmann 1986). Fis binds to two sites in the recombinational enhancer (Bruist et al. 1987) and becomes colocalized with Hin bound at each recombination site to form a three-looped invertasome structure (see Fig. 1A) that is an intermediate in inversion [Heichman and Johnson 1990]. Assembly of the invertasome may proceed by first pairing of the two recombination sites by Hin. It has been proposed that the association of the enhancer into the invertasome structure may be required to initiate rotation of the Hin subunits that are covalently attached to the core nucleotides after DNA cleavage [Heichman et al. 1991]. The nonspecific DNA-binding protein HU is believed to help stabilize the invertasome by facilitating the looping of DNA. HU strongly stimulates inversion rates only when the enhancer is located close to a recombination site, where extreme bending of the DNA is required for invertasome assembly [Johnson et al. 1986; M. Haykinson and R. Johnson, unpubl.].

Initial in vitro experiments with partially purified extracts failed to detect deletions even though inversions were readily obtained (Johnson et al. 1984). We have now reinvestigated the deletion reaction using purified proteins, different substrates, and more sensitive detection procedures. We find that site-specific deletions by the Hin system can be catalyzed in vitro, albeit at a low rate. We have used the in vitro system to determine whether the deletion reaction is mechanistically related to inversion or whether it occurs by a separate pathway, and to address the mechanism responsible for the overwhelming preference of Hin to promote inversion over deletion. We find that the deletion reaction is probably very similar to the inversion reaction and that the low rate of the deletion reaction is not a function of inefficient recognition or reactivity of recombination sites in directly repeated configuration. Rather, the directionality bias is the result of preferential assembly of recombination sites into a particular geometric configuration irrespective of starting recombination site orientation. Strand exchange is attempted and results in inversion if the sequences of the core nucleotides at synopsis are identical. However, if the core nucleotides are not identical at synopsis, as in the case of directly repeated sites, knotted products are generated. These recombinant knots can be explained by an additional DNA strand rotation that would restore the complementarity of the core nucleotides and allow

Figure 1. Pathways for assembly of recombination sites to generate inversion or deletion. In pathway A, the recombination sites are oriented in inverted [parallel] configuration with respect to each other as they are in the Salmonella chromosome. The recombination sites probably first associate on the supercoiled DNA molecule (paired hix structures) prior to assembly of the recombinational enhancer segment into the invertasome structure [Heichman and Johnson 1990]. The paired hix structures are stabilized by interactions of Hin dimers bound to each of the recombination sites. The invertasome structure is stabilized by Hin–Fis interactions and DNA supercoiling, in addition to Hin–Hin interactions. Inversion of the segment of DNA containing the enhancer occurs by double-strand cleavage followed by a 180° rotation of DNA strands and ligation in the invertasome complex [Heichman et al., this issue]. Pathways B–D represent three potential routes of assembly of recombination complexes starting with recombination sites in directly repeated orientation. Pathways B and C are two different ways on a supercoiled DNA molecule in which the recombination sites could associate with the enhancer into a configuration to generate deletion after strand exchange. If strand rotation occurs in the clockwise direction, the deletion products will be interlinked once in pathway B and unlinked [free] in pathway C. In pathway D, the recombination sites assemble into a structure that is identical to the invertasome in pathway A, except the recombination sites are aligned in reverse [antiparallel] orientation. Two rotations of DNA strands must accompany strand exchange to ligate the products [see Fig. 8]. This results in a knot being formed in the DNA but no change in the primary sequence. The results in this paper suggest that pathway B is the most likely mechanism for the inefficient deletion reaction that occurs between directly repeated recombination sites, whereas pathway D is the preferred assembly of directly repeated recombination sites.
for ligation. This results in a change in the topology of the DNA but not in the primary sequence. The results suggest further that the preferential alignment of sites within the recombination complex is directed by the topological constraints imposed by a supercoiled DNA molecule and not a molecular feature of the sites themselves. Recent studies with the Gin inversion system reach similar conclusions with regard to recombination themselves. Recent studies with the Gin inversion system reach similar conclusions with regard to recombination site synopsis [Kanaar et al. 1990].

Results

Hin-mediated deletion in vivo

The relative in vivo rates of inversion between recombination sites in inverted orientation and deletion between recombination sites in direct repeat orientation have been measured. The different plasmid substrates containing recombination sites in inverted or directly repeated orientation were transformed into a strain containing the hin gene under lacPO control [see Materials and methods]. Hin expression was induced in culture, and aliquots were removed at various times for plasmid isolation and analysis. Except for the relative orientation of their recombination sites [see Fig. 2], pMS634 [inverted repeat, IR] and pRJ858 [direct repeat, DR] are similar in structure. Figure 3 compares the rates of inversion in these two plasmids. These plasmids contain an enhancer located 868 and 694 bp from the recombination sites. Deletions occur at ~8% the rate of inversions under these in vivo conditions. In vivo recombination rates in pMS551 [IR] and pMS638 [DR] or pRJ974 [DR], which have their enhancers located at the native position 99 bp from hixL [Fig. 2], were indistinguishable from pMS634 [IR] and pRJ858 [DR], respectively [Table 1; data not shown].

Hin-mediated deletion in vitro

The preference for inversions over deletions is even greater in vitro than in vivo. Figure 4A shows the result of an in vitro Hin reaction with pRJ858 [DR]. In this experiment deletions are detected by digesting the products of the reaction with a restriction enzyme that cleaves once within the vector and radiolabeling the ends with $^{32}$P. The resulting linear parental product (5377 bp) and deletion product (3752 bp) are visualized after electrophoresis and quantitated. After 60 min, 6% of the molecules have undergone deletion. In a parallel experiment, the rates of inversion in vitro was measured in pMS634 [IR] [Fig. 4B]. We estimate that the rate of deletion in vitro is <0.3% of the rate of inversion using these substrates.

Requirements for deletion formation

To determine whether the deletion reaction requires the same reaction components as inversion, we have assayed for deletions under conditions that affect the efficiency of inversion. No deletion products were detected in reactions with pRJ858 [DR] in the absence of Fis and in substrates lacking a functional enhancer [pRJ907 [DR]; Table 2]. A requirement of DNA supercoiling for dele-

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Figure 2. Structure of the plasmid substrates used in this study. (A) The solid arrowheads denote the recombination sites, with AA or TT indicating the wild-type core sequence of hixL–WT and AT indicating the mutant core sequence present in hixL–AT. The relative orientations of the recombination sites are denoted by the direction of the arrowheads: Directly repeated recombination sites have arrowheads pointing in the same direction; inverted recombination sites have arrowheads oriented toward each other. Because of the symmetric nature of the hixC recombination sites, their relative orientation is indistinguishable. The location of the 63-bp enhancer segment is indicated by the open rectangle. pRJ907 is missing 87 bp, which includes part of the enhancer segment. The number of base pairs between the center of the recombination sites and the boundaries of the enhancer are given along with the length of the vector sequences (above the broken lines) as measured between the center of the two recombination sites. (B) Sequences of the wild-type hixL recombination site [hixL–WT], a mutant hixL recombination site [hixL–AT], and a perfectly symmetric recombination site [hixC] that is a consensus sequence of all the known DNA invertase systems related to Hin. The bold nucleotides denote the core residues where DNA strand exchange takes place.
HIN induction, aliquots were removed, plasmid DNA was prepared, and inversion or deletion was measured. The results are plotted as recombinants [inversions in pMS634 (IR), ]; or deletions in pMS858 (DR), ] per substrate molecule as a function of time after hin induction.

Deletion products are singly catenated

The requirement for the enhancer, Fis, and DNA supercoiling suggests that deletions may be generated by a mechanism similar to that used to generate inversion. Recombination intermediates that could generate deletions, assuming that the interaction between Hin and Fis at the enhancer is analogous to that proposed for inversion, are depicted in the pathways in Figure 1, B and C. Compared with the inversion complex [Fig. 1A], additional loops in DNA are required to orient the recombination sites into a configuration to generate deletion upon strand exchange. Analysis of DNA models suggests that due to the energetics of a supercoiled molecule, structure B would be more favorable than structure C. If the direction of strand rotation was exclusively in the clockwise direction, the product of recombination in structure B would be catenated with the resulting molecules interlinked once, but the deletion products generated from the intermediate depicted in structure C would not be catenated. Topological analysis of Hin-mediated recombinant knots has shown that strand rotation is exclusively in the clockwise direction in the inverted assemblage on a supercoiled molecule [Heichman et al. 1991]. A counterclockwise rotation of DNA strands in structure C would generate singly catenated products.

To determine whether the deletion products are catenated, the products of a reaction with pRJ858 (DR) were electrophoresed in an agarose gel and the DNA migrating as full-length supercoiled molecules was excised (Fig. 5A). This population will contain the parental [including knotted forms; see below] and any catenated deletion products; any free 1625-bp deletion circles will have migrated farther in the gel as denoted in Figure 5. After digestion with a restriction enzyme that cuts once

![Figure 3](image)

Figure 3. Rates of inversion vs. deletion in vivo. Expression of the *hin* gene in R12626 containing pMS634 (IR) or pMS858 (DR) was initiated by the addition of IPTG. At various times after Hin induction, aliquots were removed, plasmid DNA was prepared, and inversion or deletion was measured. The results are plotted as recombinants [inversions in pMS634 (IR), ]; or deletions in pMS858 (DR), ] per substrate molecule as a function of time after hin induction.

| Plasmid substrate | hix structure (inverted or direct repeats) | Percent inverted | Percent deleted |
|-------------------|-------------------------------------------|-----------------|----------------|
| pMS551 WT         | close                                      | 36.6            | <2             |
| pMS638 WT         | close                                      | 32.1            | 18.8           |
| pMS631 AT         | close                                      | 32.1            | 22.2           |
| pRJ857 AT         | extended                                   | 34.6            | <1             |
| pMS634 WT         | extended                                   | 34.2            | 23.1           |
| pRJ858 WT         | extended                                   | 29.7            | 21.0           |
| pRJ863 AT         | extended                                   | 36.9            | <2             |
| pRJ864 AT         | extended                                   | <1.18 bp        | 6688 bp between the enhancer and closest recombination site. |
| pMS656 hixC       | close                                      |                 |                |

\[ WT \] *hixL-WT* sites, \[ AT \] *hixL-AT* sites present on the plasmid. Their relative orientation [inverted or direct repeats] is also given.

\[ Close \] \( <1.18 \) bp, \[ extended \] \( >688 \) bp between the enhancer and closest recombination site.

\[ The percentage of inverted molecules was determined after induction of Hin for 60 min except for pMS656, which was determined after 5 hr. \]

\[ The percentage of molecules that had undergone deletion was determined 5 hr after the induction of Hin. \]
within the 1625-bp segment between the recombination sites, followed by end-labeling, the molecules were electrophoresed again and the products were detected by autoradiography [Fig. 5B]. The number of deletion products obtained was nearly identical after gel isolation to the number obtained without prior gel isolation (5.1% vs. 5.7%, respectively), indicating that the products must be catenated.

Electron microscopy of deletion products that were coated with RecA was performed to determine the complexity of catenation [Krasnow et al. 1983]. For this experiment we used pRJ864, which contains two hixL–AT sites in direct repeat orientation. This plasmid supports wild-type rates of deletion [see below] but does not knot extensively, which simplified the search for deletion products [Heichman et al., this issue; data not shown]. To enrich for deletions, the products of a reaction with pRJ864 (DR) were nicked with DNase I and electrophoresed in an agarose gel. Aliquots were removed at the times indicated, digested with NdeI, which cuts once in the vector sequences, end-labeled with $^{32}$P, electrophoresed in an agarose gel, and autoradiographed. The arrow indicates the labeled deletion product. [B] The amount of deletion products from A was quantitated and plotted as deletion products (■) per substrate molecule as a function of time. For comparison, the number of inversions (●) per molecule in pMS634 (IR) as a function of time from a parallel experiment is shown.

**Figure 5.** Hin-mediated deletion products are catenated. pRJ858 (DR) was reacted with Hin, Fis, and HU for 60 min and electrophoresed in an agarose gel (A, lane 2) together with unreacted supercoiled DNA (lane 1). The location of a potential free 1625-bp supercoiled deletion product is indicated as determined by electrophoresing supercoiled plasmids of known size on the same gel. The DNA migrating in the region of the parental supercoiled (SC) and knotted forms, as indicated, was isolated, cleaved with HindIII, which cuts within the 1625-bp segment between the recombination sites, end-labeled with $^{32}$P, electrophoresed on a second agarose gel, and autoradiographed (B, lane 3). Lane 1 (B) is HindIII-cut end-labeled pRJ858 (DR) that was not reacted with Hin; lane 2 is an aliquot of the same reaction used in A that was digested with HindIII and end-labeled without gel isolation. The location of the 1625-bp labeled deletion product, as well as the parental DNA, is indicated.

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**Table 2. Requirements for Hin-mediated site-specific deletion**

| Reaction | Substrate | Relative number of deletions |
|----------|-----------|-----------------------------|
| A        |           |                             |
| complete | pRJ858    | 1.0                         |
| no Fis   | pRJ858    | <0.02                       |
| no enhancer | pRJ907 | <0.02                       |
| no supercoiling | pRJ858 | <0.02                       |
| B        |           |                             |
| complete | pRJ858    | 1.0                         |
| no HU    | pRJ858    | 0.57                        |
| 3× HU    | pRJ858    | 0.73                        |
| complete | pMS638    | 0.17                        |
| no HU    | pMS638    | 0.02                        |
| 3× HU    | pMS638    | 0.34                        |

*Reactions were performed for 60 min at 37°C in the presence of 70 ng of Hin, 25 ng of Fis, and 100 ng of HU on 0.1 pmole of supercoiled plasmid substrate (complete) unless denoted otherwise. [3× HU] 300 ng of HU was included in the reaction. The percent of plasmids containing deletions are presented relative to pRJ858 in complete reaction conditions. In experiment A, pRJ858 gave 5.2% deletions; in experiment B, pRJ858 gave 8.6% deletions. pRJ858 was relaxed with topoisomerase I prior to reaction.
resed in an agarose gel. The DNA that migrated faster than parental relaxed molecules was purified, coated with RecA, and examined with the electron microscope. Two catenated deletion molecules were found, and each was interlinked once [Fig. 6]. Ten deletion products generated with another plasmid that supports deletion [pMS631 (IR); see below] were also determined to be exclusively singly catenated.

Substrates containing wild-type recombination sites in direct repeat orientation support efficient knotting

In the work of Heichman et al. (this issue), it has been shown that the knotting of DNA by Hin is mechanistically related to recombination and is caused by multiple rotations of cleaved DNA strands prior to ligation. Although it only supports a low rate of deletion, pRJ858 (DR) is a very efficient substrate for Hin-mediated knotting. Figure 7 [left] shows the result of a Hin reaction time course in which pRJ858 (DR) was nicked with DNase I to remove supercoils prior to electrophoresis. A series of knots containing from 3 to >15 nodes is generated with increasing time. The primary knotted products are multiples of three, giving rise to 3-, 6-, 9-, 12-, and 15-noded molecules. In addition, an increasing number of molecules with 5, 8, 11, and 14 nodes are also formed. The pattern of knots in pRJ858 (DR) corresponds precisely to those generated in pRJ862 (IR), which contains a mutation within the core nucleotides of one recombination site [Fig. 7 (right); Heichman et al., this issue]. This demonstrates that substrates with directly repeated recombination sites and substrates containing inverted recombination sites with core mutations are equally as reactive for knotting. These results lead to a model in which plasmids containing directly repeated sites are efficiently forming a recombination complex that is identical to that formed normally between inverted sites [see Figs. 1D and 8B]. In this assembly, a single 180° rotation will position the core nucleotides into a mismatched configuration. Therefore, an additional rotation leading to the formation of a three-noded knot would be required to achieve ligation once strand exchange has been initiated. Two additional rotations will generate a five-noded product, and multiple independent reactions will lead to more complex knots [Heichman et al., this issue].

Recombination between substrates containing symmetric core nucleotides

The model described above suggests that plasmids containing their recombination sites in directly repeated orientation assemble efficiently into an invertasome complex in which the recombination sites are in a reverse or antiparallel orientation. The inability to invert the intervening DNA between directly repeated sites is primarily or exclusively a function of the nonidentity of the core nucleotides, which prevent ligation after a 180° rotation. This model was tested with recombination sites containing a symmetric 2-bp core sequence. In this case, ligation after a single 180° rotation of strands could be achieved regardless of the starting arrangement of recombination sites at synapsis [see Fig. 8C]. Thus, a plasmid with two recombination sites containing symmetric core sequences in directly repeated configuration should support a high rate of inversion, as well as a low rate of deletion.

Plasmids were constructed with both recombination sites containing symmetric 2-bp core sequences.
sites containing AT for their core nucleotides [hixL-AT]. The recombination sites were positioned in either inverted or directly repeated orientation, and the enhancer was placed either close to one recombination site [pMS631 (IR) and pRJ857 (DR)] or well spaced from both recombination sites [pRJ863 (IR) and pRJ864 (DR)] (Fig. 2). Each of these plasmids was assayed in vitro and in vivo for their ability to support inversion or deletion. High rates of inversion, which were similar to those in plasmids containing hixL-WT sites in inverted orientation, were measured for the plasmids containing hixL-AT sites in either orientation (Fig. 9; Table 1; data not shown). Each of the hixL-AT plasmids also supported a low rate of deletion that was similar to plasmids containing hixL-WT sites in direct repeat (Fig. 10; Table 1; data not shown). No deletion or inversion products were observed in plasmids containing hixL-WT sites in inverted or direct repeat orientation, respectively. These results clearly demonstrate the importance of the core nucleotides in determining the product of recombination but indicate that the relative orientation of the recombination sites is not important for reactivity.

The apparent preference for assembly of recombination sites into the invertasome structure, regardless of the sequence of the recombination sites, was investigated further by assaying the product of recombination with a plasmid containing hix sites that are symmetric over their entire sequence. pMS656 has recombination sites with a palindromic sequence that is a consensus of the DNA invertase family [hixC; see Fig. 2B]. Because the recombination sites are completely symmetric, their relative orientation cannot be distinguished. Although pMS656 does not recombine as efficiently as the plasmids with hixL-AT sites, a >10-fold bias of inversion over deletion is measured in vitro (data not shown) and in vivo (Table 2). These results argue further that the nucleotide sequence of the recombination site outside of the core sequence is not responsible for mediating directionality.

Discussion

We have characterized the Hin-promoted deletion reaction that excises DNA between two hix sites. The deletion reaction is much less efficient than inversion. In the presence of moderately overproduced Hin protein, the rate of deletion in vivo is <10% of the rate of inversion in plasmids with appropriately oriented recombination sites. The difference in the relative rates of the two reactions is even more pronounced in vitro, where deletions are produced at 0.3% of the rate measured for inversion. In vitro experiments have shown that the deletion reaction requires DNA supercoiling, the enhancer DNA segment, and Fis, suggesting that the nucleoprotein complex that is assembled to promote deletion is similar to the invertasome structure that is the precursor of inversion. The most significant difference is the requirement for additional loops of DNA to orient directly repeated recombination sites into a configuration that will yield deletion. On the basis of topological constraints imposed by supercoiling and the finding that the deletion products are singly catenated, the loops are likely to be positioned as depicted in Figure 1B.

Determinants of recombination site alignment

Does the low rate of deletion reflect the inefficiency of two sites in directly repeated configuration to be effi-
Figure 9. Rates of in vitro inversion of wild-type and mutant substrates. pMS634 [IR] (●), pRJ863 [IR] (□), pRJ864 [DR] (■), and pRJ858 [DR] (▲) were incubated with Hin, Fis, and HU for various times, digested with restriction enzymes, and electrophoresed in an agarose gel to determine the fraction of molecules containing inversions. The results are plotted as inversions per substrate molecule as a function of time. pMS634 [IR] and pRJ858 [DR] contain two hixL-WT sites, and pRJ863 and pRJ864 contain two hixL-AT sites.

It was surprising to find that a large length of DNA past each other would preferentially generate collisions between sites in a configuration that would lead to inversion upon strand exchange (Benjamin and Cozzarelli 1986). The association of sites may occur in a branched DNA molecule to directly give the invertasome structure that contains the enhancer, or more likely, the two recombination sites may first assemble into a complex prior to association with the enhancer. Both types of complexes have been observed by electron microscopy, and their formation has been shown to be strongly stimulated or dependent on DNA supercoiling (Hiechman and Johnson 1990). For recombination sites in inverted orientation, the favored (parallel) alignment of sites will generate inversion products upon a single 180° rotation (Figs. 1A and 8A). For recombination sites in direct repeat orientation, the sites will join in an antiparallel alignment as in pathway D of Figure 1. A single 180° rotation of strands results in core sequences that are unable to base-pair (Fig. 8B). A second rotation is required to allow for base-pairing and ligation. This leaves the primary structure of the DNA unchanged; however, because of the geometry of the DNA strands in the invertasome, a knot is generated. Alignment of the recombination sites into a configuration that will lead to singly catenated deletion products necessitates that one of the sites be turned 180°, generating the structures depicted in Figure 1, pathway B. The assembly of these structures would not be kinetically favored on supercoiled DNA, and the structures themselves would be under severe torsional stress that would thermodynamically discourage the same interwrapping of sites found in the inversion complexes.

The overwhelming propensity for assembly of the Hin recombination sites into a configuration that leads to inversion may be due exclusively to constraints placed by the conformation of the supercoiled DNA substrate. Two mechanisms may be contributing to the biased assembly of recombination sites; these mechanisms are not necessarily mutually exclusive. The first is that a particular juxtaposition or interwrapping of protein-bound DNA sites is favored on supercoiled DNA (Craigie and Mizuuchi 1986; Boocock et al. 1987; Gellert and Nash 1987; Stark et al. 1989). This interwrapping may be required to successfully assemble a complex competent to promote recombination and will not favor formation of a deletion complex. The second is that plectonemically wound DNA strands that are continually sliding efficiently recognized or assembled into a reactive complex? This cannot be the case for several reasons. A plasmid substrate containing the wild-type recombination sites in directly repeated configuration supports an efficient knotting reaction with rates similar to that observed with a substrate containing inverted sites with nonhomologous core residues. Most importantly, plasmids containing directly repeated sites with symmetric core nucleotides undergo an efficient inversion reaction, as well as an inefficient deletion reaction. The combined results indicate that recombination sites in directly repeated orientation are efficiently assembled into an invertasome complex that is capable of initiating strand exchange.

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greater length. However, even with one large segment of DNA between the enhancer and a recombination site, very low deletion rates were observed if the other segment was at the wild-type spacing. It is possible that stress placed on the deletion intermediate structure by the additional looping of DNA may distort the structure sufficiently to inhibit association of the enhancer with the second recombination site separated by a short DNA segment. The effect of DNA length on deletion rates can be partially overcome by additional HU. In vivo, the requirement for large segments of DNA between both recombination sites and the enhancer is not observed. The interaction of DNA in vivo with chromosomal binding proteins like HU, perhaps together with the reduced superhelical tension of DNA in the cell (Sinden et al. 1983; Bliska and Cozzarelli 1987), may relieve some of the topological stress in the deletion complex. These factors may also contribute to the reduced preference of inversions over deletions measured in vivo as compared with in vitro.

We conclude that the product of Hin-mediated recombination is governed by two factors. The first is the choice of assembly of recombination sites into a nucleoprotein complex to yield inversion or deletion of the intervening DNA. Probably because of the nature of supercoiled DNA, the sites preferentially assemble into a configuration that yields inversion. Once a recombination complex is assembled, the sequence of the core residues determines whether an attempt at inversion or deletion will be successful. Ability of the strands to ligate following an initial 180° rotation results in the formation of recombinants. If the DNA strands cannot be ligated after a single 180° rotation of strands, further rotations in the same direction can occur to achieve ligation; however, this results in the formation of knots in the DNA. A similar mechanism for a reaction between directly repeated sites has been proposed independently for the Gin inversion system (Kanaar et al. 1990).

Recent analysis of another related site-specific recombination system, resolution by Tn3 resolvase, has provided valuable insight into recombination site alignment. The resolvase system displays a strong preference for a particular configuration of sites that gives rise to resolution over inversion (Reed and Grindley 1981; Krasnow and Cozzarelli 1983), just as the invertases show such a strong preference for inversion over deletion. Unlike Hin, however, resolvase does not require any accessory proteins such as Fis, nor does it require a recombinational enhancer. The res recombination sites are more complicated than hix sites, containing three resolvase protein-binding subsites (Grindley et al. 1982). These additional binding sites have been shown to dictate the strict specificity of resolvase-mediated recombination [Dröge and Cozzarelli 1989; Stark et al. 1989]. Thus, like the Hin system, synopsis of the res recombination sites is not determined exclusively by a particular parallel or antiparallel orientation [Bednarz et al. 1990]. Rather, alignment is dependent on the preferential formation of a particular synaptic intermediate, which leads to the bias of one recombinant product over the other.

Implications for in vivo recombination by Hin

The structure of the hix recombination site is relatively simple, particularly in comparison with other site-specific recombination sites utilized in systems such as λ and resolvase. It has been observed that Hin can bind to a variety of pseudo-hix sites (Glasgow et al. 1989) and, in some cases, can recruit these sites into a recombination complex that can initiate cleavage [Johnson and Bruist 1989; Heichman and Johnson 1990]. The only criterion left for the system to determine whether the chosen sites will recombine to change the primary structure of the DNA is the identity of the 2-bp core nucleotides. Given random sequence, recombination will be productive 1 of every 16 times. The remainder of the events will undergo an additional rotation to restore the original sequence. This may lead to an unacceptably high rate of Hin-promoted chromosomal rearrangements. However, several additional levels of control operate to limit unwanted rearrangements. Hin protein is synthesized at an extremely low level, and it has been shown that the amount of Hin limits the rate of the inversion reaction controlling H2 flagellin expression [Bruist and Simon 1984]. In addition, the Hin protein cannot efficiently catalyze inversion or deletion without the presence of a recombinational enhancer sequence located somewhat nearby. The maximum distance allowed for function of the recombinational enhancer is not known, but the observation that inversion rates are reduced by threefold in vitro when the enhancer is 4 kb from the closest hix site suggests that it may not function over extremely great distances [Johnson et al. 1986]. The constraints on the structure of the recombinational enhancer (two Fis-binding sites separated by 48 or 58 bp) would limit the number of these sites present in the chromosome [Johnson et al. 1987]. Finally, the potential biological consequences of a deletion (the less favored reaction) would be far greater than an inversion, which will only cause mutations at the site of recombination on the chromosome.

Materials and methods

Enzymes and reagents

Purified Hin, Fis, and HU preparations were obtained as described in Heichman et al. [this issue]. Restriction enzymes and isopropylthio-β-D-galactoside (IPTG) were purchased from Boehringer Mannheim or New England Biolabs; AMV reverse transcriptase was from Promega; and calf thymus topoisomerase I was from Bethesda Research Laboratories, Life Technologies, Inc. Antibiotics were obtained from Sigma.

In vitro Hin recombination reactions

The reaction conditions used to generate inversions, deletions, and knots were identical except where noted and are described (Heichman et al., this issue). Five percent dimethylsulfoxide was included in reactions with pMS656. Inversions were detected by digestion of the reacted plasmids with the appropriate restriction enzymes to determine the orientation of the invertible segment followed by agarose gel electrophoresis. Deletion products were typically detected as follows: Reacted plasmids were digested with a restriction enzyme that cleaves the DNA.
once within the vector or deletion segment and then end-filled with 10 μCi [α-32P]dATP (Amersham, >3000 Ci/m mole) with AMV reverse transcriptase. The labeled DNA molecules were detected by autoradiography after agarose gel electrophoresis. The reactions were quantitated by Cerenkov counting of gel with 10 IzCi [α-32P]dATP (Amersham, >3000 Ci/mmole) with low. pMS551, pMS631, and pMS634 have been described previously [Johnson and Simon 1985]. pBR322, have one recombination site located at the ori, tet

In vivo Hin recombination reactions

In vivo recombination reactions were performed in RJ2626 [pRJ918, pRJ823, Δlac-pro] tecA56 rpsL ara srl]. This strain contains the hin gene tightly regulated by the lac promoter on pRJ918 such that under standard conditions, no inversion or deletion is detectable until IPTG is added. Substrate plasmids were transformed into RJ2626 and plated on LB agar containing 100 μg/ml of ampicillin, 15 μg/ml of tetracycline, 50 μg/ml of spectinomycin, and 0.2% glucose. Single colonies appearing after overnight growth were inoculated into LB plus antibiotics and grown at 37°C until OD600 = 0.3 at which time IPTG was added to 1 mm. Aliquots were taken at appropriate times, and plasmid DNA was isolated by the alkaline–SDS lysis method [Maniatis et al. 1982]. Cultures were maintained in exponential phase by continual dilution with prewarmed media. Inversions or deletions were detected as described above.

Plasmid structures and constructions

The overall structure of each of the substrate plasmids used in this study is similar. The plasmids, which are all derivatives of pBR322, have one recombination site located at the EcoRI site and one recombination site located at the Sall site. The relative orientations of the two recombination sites and the locations of the enhancer segment are depicted in Figure 2. The details of the structures and constructions of the plasmids are described below. pMS551, pMS631, and pMS634 have been described previously [Johnson and Simon 1985]. pMS638 and pRJ857 are identical to pMS551 and pMS631, respectively, except that the hixL sequence at the Sall site is oriented in directly repeated orientation with respect to the hixL at the EcoRI site. pRJ858 was constructed by a ligation of the EcoRI-HindIII DNA fragment from pMS634, which contains hixL plus the enhancer with the EcoRI-HindIII fragment from pMS638, which contains the pBR322 ori and hixL. pRJ907 was made from pRJ858 by deleting the 87-bp segment of DNA between the Clal site in the enhancer and the unique HindIII site. This removes the distal domain and most of the intervening DNA between the two domains of the enhancer. The construction of pRJ862 is described in Heichman et al. [this issue]. pRJ863 and pRJ864 are derived from pMS631 and pRJ857, respectively, by inserting into the Smal sites of these plasmids a 789-bp HindIII fragment containing the lacPL162 mutation [Reznikoff et al. 1982]. The same 789-bp fragment was ligated into an end-filled HindIII site in pMS638 to give pRJ974.

pMS556 was constructed as follows: The oligonucleotide d(TATCATCAATAACATGTTAGGATAA), encoding a consensus and completely symmetric recombination site (hixC), was ligated into the Smal site of M13mp8 to give mMS500. The EcoRI–BamHI fragment containing hixC was ligated into similarly cut pMS577 [Johnson and Simon 1985], which contains the enhancer, to give pMS555. The EcoRI–SalI fragment containing hixC from mMS500 was also ligated into the Sall site of pBR322 with T4 DNA polymerase to fill in the noncomplementarity ends to give pMS653. pMS656 was constructed from these two plasmids by substituting the EcoRI–HindIII fragment from pMS655, which contains hixC plus the enhancer for the 29-bp EcoRI–HindIII region in pMS653.

pRJ918 [lacP, p15A ori, tet] was generated by transferring the EcoRI fragment containing the lacP gene from pMS421 [M. Susskind, University of Southern California] into the EcoRI site of pACYC184 [Chang and Cohen 1978]. pRJ918 contains the lacPO region fused to the hin gene 40 bp before the initiating AUG in a low-copy-number replicon. It was constructed by ligating a PeuII–HindIII fragment containing lacPO-hin from pMS568 [Johnson and Simon 1985] into the Smal–HindIII linker region of pMS421 [lacP, pSC101 ori, spc/st]

Electron microscopy of deletion products

RecA coating and electron microscopy of DNA molecules was performed as described in Heichman et al. [this issue]. pRJ864 or pMS631 was reacted for 30 min and electrophoresed through agarose gels after nicking with Dnase I. The DNA that migrated faster than the parental nicked species was excised, purified with the Gene clean protocol [Bio 101], and coated with RecA protein.

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