Retroviral DNA Integration

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INTRODUCTION

Integration is an obligatory replication step for all retroviruses. The process begins after the virus enters the cell and the RNA genome is reverse transcribed into a double-stranded DNA. The direct precursor to integration is the linear form of viral DNA (4, 64). Circular DNA forms are detected in cells but are considered to be dead-end reverse transcription products (49) or products resulting from autointegration events (63). Linear viral DNA contains at its termini long terminal repeats (LTR) sequences. The ends of these LTRs are specifically recognized by the viral integrase (IN). Preintegration complexes, capable of catalyzing integration in vitro, can be isolated from infected cells (30). These complexes contain linear viral DNA, several viral proteins including matrix (7), reverse transcriptase (63), nucleocapsid (59), and IN (30) and at least two cellular proteins, high-mobility-group [HMG-I(Y)] (29) and barrier to autointegration factor (BAF) (62). Depending upon the retrovirus, preintegration complexes either enter the nuclei of nondividing cells through the nuclear pore (e.g., human immunodeficiency virus [HIV]) or wait until the nuclear membrane dissolves during cell division (e.g., Moloney murine leukemia virus [MoMuLV]) (75). In some retroviruses, nuclear localization signals are associated with various viral proteins, including IN, and facilitate migration to the nucleus (34, 55, 84). Once the preintegration complex associates with the host chromosome, viral IN catalyzes the insertion of the viral sequences into the host DNA. The two LTR ends of the linear viral DNA are brought together into a ternary complex with IN and host DNA, where the insertion occurs in a coordinated or concerted reaction. A 2-bp sequence is lost from each end of the viral DNA, and a short duplication of 4 to 7 bp from the host, depending upon the viral IN, is introduced into the host sequence flanking the viral DNA. Repair of damage to the host DNA by integration is presumably mediated by cellular enzymes. Integrated viral DNA is termed the provirus.

IN was first detected in avian myeloblastosis virus as a non-specific endonuclease of 32 kDa. Sequences from peptide fragments of the protein showed sequence homology to the β chain of avian sarcoma virus (ASV) reverse transcriptase (77). An active enzyme consists of a multimeric structure of at least a dimer (47). The primary evidence establishing that the endonuclease was encoded in the viral genome came from analysis of an IN mutant ASV, LA335, which was temperature sensitive for replication and possessed a temperature-sensitive DNA endonuclease (36). These observations were confirmed by experiments in which mutations were introduced into conserved residues of IN and blocked replication of the virus (73). The first biochemical evidence linking the DNA endonuclease to integration was the demonstration that the enzyme could specifically cleave viral LTR ends (19). It was subsequently shown that IN could also preferentially bind to the LTR termini of viral DNA (54, 69). However, it was not until DNA oligodeoxynucleotides representing the ends of the viral LTRs were used as substrates for integration in vitro that IN was shown to be both necessary and sufficient to mediate integration (51) and that an energy source such as ATP was not required for the reaction.

STRUCTURE OF INTEGRASE

IN, encoded in the pol gene of the virus, is translated as part of a large Gag-Pol polyprotein and is processed into its mature form by the virus-encoded protease (PR). Based upon evidence from limited protease digestion studies (24) and alignment of the primary sequences of several IN proteins (46), which identified clusters of conserved residues, IN is thought to
possess three structural domains. The domains consist of an N-terminal domain of 50 amino acids with a putative zinc binding motif resembling a zinc finger (HHCC), a central domain of 160 amino acids with a D,D(E)35 motif, and a less highly conserved C-terminal domain of 80 amino acids. The crystal structures of the central ASV and HIV-1 IN core domains have been solved (6, 20). Nuclear magnetic resonance spectroscopy (NMR) solution structural data is available for the HIV-1 IN N and C-terminal domains (11, 12, 21, 22, 65). Figure 1 displays each of the known fragment structures of HIV-1 IN. Unfortunately, there is no structural data for an intact protein with or without a substrate mimic, so that we do not know how these fragments fit together in the holoenzyme.

N-Terminal Domain

Within the N-terminal domain of IN is a putative zinc binding motif resembling a zinc finger (HHCC), a central domain of 160 amino acids with a D,D(E)35 motif, and a less highly conserved C-terminal domain of 80 amino acids. The crystal structures of the central ASV and HIV-1 IN core domains have been solved (6, 20). Nuclear magnetic resonance spectroscopy (NMR) solution structural data is available for the HIV-1 IN N and C-terminal domains (11, 12, 21, 22, 65). Figure 1 displays each of the known fragment structures of HIV-1 IN. Unfortunately, there is no structural data for an intact protein with or without a substrate mimetic, so that we do not know how these fragments fit together in the holoenzyme.

Central Core Domain

The central core domain comprises residues 50 to 235 and has been shown to coordinate divalent cations. The crystal structures of the catalytic core domains for HIV-1 (20) and ASV (6) have been solved. Crystallization of the HIV-1 core was dependent upon a F185K substitution, which increased its solubility (20, 44, 45); this is shown in Fig. 1. While an intact IN protein is required for complete activity, an ASV fragment is capable of end processing but not joining. The HIV-1 core fragment is not capable of either reaction (6, 40). Both cores, however, catalyze the disintegration reaction (6, 40, 45). The overall folded structure of the catalytic core is similar to that of nucleases such as Escherichia coli RNase H, the HIV-1 RNase H domain of reverse transcriptase and E. coli RuvC (6, 74, 88).

C-Terminal Domain

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dinucleotide in the viral LTR resides in the catalytic core, implying that the core contains the active site (52). The core domain contains a catalytic triad of three highly conserved residues, D,D(35)E. The triad for ASV is D64, D121, and E157. Substitutions of any of these residues generally abolish end-processing and/or joining reactions (57). The comparable triad for HIV-1 IN is D64, D116, and E152 (Fig. 1). Here again, most substitutions inactivate the enzyme, with two exceptions: a D116N substitution results in a more active HIV-1 IN than wild type in vivo (25), and an enzyme with D116E and E152D substitutions, while incapable of end processing and joining, catalyzes detectable disintegration reactions (24). Crystal structures of the catalytic core, coordinating a divalent cation, have been determined for ASV by using Mg\(^{2+}\) and Mn\(^{2+}\) (5) and for HIV by using Mg\(^{2+}\) (35, 68). The divalent cations were found to be coordinated by the two conserved aspartic acid residues of the catalytic triad. A comparison of HIV-1 and ASV cores indicated that the two aspartic acid residues are similarly positioned within the respective HIV-1 and ASV structures (5, 6, 20, 35, 36, 68). A relatively extensive mutational analysis of conserved residues in the catalytic core of the intact IN has been performed (18, 20, 24, 44, 45, 53, 57, 58, 60). While some of the substitutions affect end-processing, joining, and disintegration reactions to various extents, others are less disruptive and in some cases allow end processing but prevent joining. These results imply that not all conserved residues in the core are essential to catalytic activity.

C-Terminal Domain

The C terminus of IN is the least highly conserved of the three domains (13, 67). An HIV-1 fragment representing residues 235 to 288 binds nonspecifically to DNA (26, 53, 66, 71, 82, 87). The structure of the C-terminal domain of HIV-1 IN, residues 220 to 270, has been determined by NMR (21, 65) and contains an SH3 fold motif (Fig. 1). Such structures have been found in a number of proteins but only in one other protein involved in DNA binding, Sso7d (3, 21, 65). In addition, multimerization of IN appears to be defective, suggesting that the C terminus of the protein is needed for dimerization or multimerization of the enzyme. Nuclear localization signals, which facilitate the entry of preintegration complexes into the nucleus, have been mapped to the C-terminal IN domain of several retroviruses and transposons. In ASV IN, the nuclear localization signal contains both basic and proline residues (55).

MULTIMERIZATION OF INTEGRASE

Several groups have examined the parameters of multimerization both in vitro and in vivo (23, 32, 80). While truncated N- or C-terminal HIV-1 IN fragments are incapable of supporting end-processing or joining reactions, a mixture of N-terminally truncated IN and C-terminally truncated IN has detectable activity (23, 80). These results imply that HIV-1 integrase is active as a dimer or higher-order multimer and that only a single N- or C-terminal domain is required for activity. In other mixing experiments, IN proteins containing single substitutions in the catalytic triad of the core domain could be complemented by IN proteins containing truncations to either the N- or C-terminal domains (23, 80). By using IN monomers having truncations in the N or C terminus, cis to the catalytic core and complemented with monomers having the missing truncated region, it was shown that the C-terminal domains cis to the catalytic site were active in end-processing and joining reactions in vivo whereas N-terminal truncations were not active in cis (23, 80).

MECHANISM OF RETROVIRAL INTEGRATION

Integration occurs in two well-characterized catalytic steps, referred to as end processing and joining, respectively. End processing involves removal of a dinucleotide, adjacent to a highly conserved CA dinucleotide, from the 3’ strand of the U3 and U5 viral DNA LTRs in a reaction involving a water molecule or other nucleophile (27, 83) (Fig. 2). This exposes a 3’ hydroxyl group, whose oxygen is used as an attacking nucleophile on the target DNA during the joining reaction, in which the viral DNA is inserted into the cellular DNA (27, 83). Engleman et al. identified three different forms of the cleaved dinucleotide resulting from end processing, depending upon the nucleophile used in the reaction (27). The most abundant form was a dinucleotide with a 5’ phosphate and a 3’ hydroxyl, which would arise when a water molecule acted as the nucleophile. It is believed that a Mg\(^{2+}\) atom coordinated [through the conserved D,D(35)E residues] in the active site of IN facilitates the deprotonation of the water to activate it as a nucleophile. This mechanism is analogous to the polymerization reaction catalyzed by DNA polymerases, including reverse transcriptase, or to activity catalyzed by adenosine cyclase (48, 79, 89), although in this case IN is believed to coordinate one rather than two Mg\(^{2+}\) atoms. A second dinucleotide product was detected when gliceral, introduced together with IN, acted as the nucleophile. This species was poorly phosphorylated and was resistant to phosphatase activity, implying the absence of a 5’ phosphate (27, 83). The third dinucleotide was identified as a 3’-5’ cyclic pGpT\(_{\text{OH}}\). This form arises when the hydroxyl of the terminal T acts as the attacking nucleophile to break a phosphodiester bond on the same strand (27).

The actual mechanism of viral DNA insertion was shown, by use of a phosphothioate-substituted target, to invert the chirality of the phosphate at the site of insertion (27). This indicates that a single-step transesterification reaction occurs and eliminates the possibility that IN forms a covalent intermediate with DNA. Such enzyme-DNA intermediates are found for topoisomerases (14) and for IN of lambda bacteriophage (70, 81), where the chirality of the electrophile phosphate is maintained in joining the DNA.

The IN recognition sequence within the LTRs is relatively short. MuLV IN recognizes 11 to 12 bp (76), ASV IN recognizes 15 bp (51), and HIV-1 IN recognizes 20 bp (78). Cross-linking and substitution of bases in the LTR of HIV-1 have demonstrated that specific interactions between IN and the terminal LTR sequences are required for end-processing and joining reactions (28, 38, 39). The sequences at the U5 and U3 LTR ends are both derived from nearly perfect inverted repeats. Addition or deletion of sequences at or near the conserved CA dinucleotide at the termini of the LTRs alters the efficiency of DNA integration (17). For example, addition of sequence 3’ to the conserved CA dinucleotide in MuLV LTRs resulted in a delay in growth of the virus by several days. However, subsequent passages of the mutant virus selected for viruses with wild-type growth rates and concomitant nucleotide changes in the LTR sequences. One revertant contained a C-to-A transition 4 bases from the 3’ termini; another contained an 11-base deletion such that integration utilized a new internal CA dinucleotide. Replacement of 2 bases in the ASV U5 LTR adjacent to the CA dinucleotide or 4 bases adjacent to and including the C of the CA dinucleotide also decreased the
efficiency of integration and resulted in a delayed-growth phenotype (16).

EARLY IN VITRO-RECONSTITUTED INTEGRATION SYSTEMS

Fujiwara and Craigie (33) described the first cell-free integration system that used extracts of MoMuLV, a large linear donor DNA containing modified LTR termini with 5'-9 overhangs, and lambda DNA as the target. The donor DNA was constructed by placing an NdeI cleavage site between tandemly linked U5 and U3 LTR termini inserted in a plasmid. After restriction digestion, a linear donor was produced with LTR termini that differed from wild-type LTR termini by 1 bp and contained 5'-2-base AT overhangs rather than blunt ends. Katz et al. (50) used purified ASV IN to reconstitute integration with a similar NdeI-constructed ASV donor DNA. Integrants in both systems were analyzed by packaging the lambda acceptor DNA, introducing the phage into cells, plaque purifying, and sequencing. Integrants displayed properties characteristic of in vivo retroviral integration, including the loss of 2 bp from the ends of both LTR termini, short duplications of the lambda DNA at the site of integration, and random distribution of integration sites.

CONCERTED DNA INTEGRATION

Murphy and Goff (72) demonstrated that when deletions were placed in the U3 LTR, 5' to the conserved CA dinucleotide, end processing in both U3 and U5 LTRs was adversely affected. This result implies that integration in vivo occurs by a concerted mechanism in which the two LTR ends of the viral DNA are inserted into a single target site, such that a mutation in one LTR can influence the processing of the other. Unfortunately, oligodeoxynucleotide model substrates do not exhibit the concerted properties characteristic of in vivo integration. One exception was described by Kukolj and Skalka (56), who designed short duplex substrates whose sequences matched those of U3 and U5 ends of ASV and HIV-1 DNA but were covalently synapsed across the termini by short single-stranded linkers. These substrates were used more efficiently than were unlinked oligodeoxynucleotides duplexes. Moreover, substrates with a paired wild-type and mutated terminus were cleaved poorly at both ends, indicating that when termini were juxtaposed, the processing of both ends displayed concerted behavior. By using tethered donor molecules, the optimum spacing for the ASV system was shown to be 2 nucleotides. If both strands were tethered, the efficiency of the reaction was considerably decreased, due in part to a loss of torsional flexibility imparted by the gap in one strand. These results provided biochemical evidence that molecular communication must take place between IN bound to both viral DNA ends.

RECONSTITUTION OF AVIAN SARCOMA VIRUS CONCERTED DNA INTEGRATION

A concerted integration system using purified IN was reported by Fitzgerald et al. (31). The system included purified...
AMV IN; lambda as the linear acceptor DNA; a donor DNA, 3.4 kb in length, containing 30 bp from the ends of the LTRs with preprocessed 5’ AT overhangs; and a supF suppressor gene as a selectable marker. The presence of the supF RNA gene in the mini-donor DNA provides for genetic selection of individual integrants in bacterial cells containing an expression vector with antibiotic resistance genes with amber mutations in the coding sequences. The linear donor was again constructed by using the NdeI restriction enzyme site (31). A small percentage of the integrants isolated from the reaction exhibited concerted integration and displayed properties expected of ASV DNA integration in vivo. However, the remainder arose through nonconcerted integration events, which produced deletions in the acceptor DNA. Vora et al. (86) established a similar system by using a 487-bp donor DNA that was also with precleaved NdeI ends. Here again, products arose from both concerted and nonconcerted reactions. Further analysis of the concerted integration products showed that they resulted from two one-ended integration events by different donor DNAs into the same acceptor rather than from both ends being provided by the same donor. As a consequence, U3-U5, U3-U3, or U5-U5 donor combinations were detected, complicating the analysis of the integration product (31, 86). In subsequent reports, changing of the buffer conditions (31, 85, 86) improved the overall efficiency of integration but with different percentages of concerted products detected.

The ASV reconstituted system that appears to most closely approximate the concerted integration in vivo uses purified recombinant ASV IN, a 3.4-kb supercoiled or linear target DNA, and a “mini” linear donor DNA substrate of only 294 bp with blunt authentic unprocessed viral termini (1, 42). This small size was chosen to maximize the probability that ends from the same donor would come in contact to facilitate concerted integration. The donor contains a supF suppressor tRNA gene flanked by only 15 bp from the ASV LTR termini. To increase the integration efficiency of this system, a host cell protein was also added to the reaction mixture by Aiyar et al. (1). The host cell protein was from the HMG family and has the ability to bend DNA that could assist in juxtaposing the U3 and U5 LTR termini in an integration complex. This bending of the donor DNA could favor concerted DNA integration. While only a small percentage of the donor DNA was integrated into the acceptor in this system, more than 90% of the integrants detected used a concerted mechanism with a single donor molecule (1). Removal of either the U3 or the U5 LTR sequences from the donor substrate resulted in a substantial reduction in the total number of integration products detected. HMG-1 was the initial HMG protein family member added to this system. It stimulated the integration reaction about fourfold compared with the activity observed in the presence of IN alone (1). Subsequently, Farnet and Bushman (29) reported that an HMG protein family member, HMG-I(Y), could be detected in HIV-1 preintegration complexes isolated from infected-cells and that integration was dependent upon the continued presence of HMG-I(Y). The addition of HMG-I(Y) to the ASV mini-donor DNA reconstituted system stimulated integration by more than 10-fold, with the mini-donor DNA being integrated via a concerted mechanism during the course of the reaction (42). Individual integrants, isolated from reactions reconstituted in the presence of HMG-1 or HMG-I(Y), showed end processing, concerted insertion with base pair duplication of acceptor DNA flanking the integrated donor DNA, and non-sequence-specific integration of the donor into the target, all characteristic of in vivo integration (1, 42).

**MUTATIONS IN THE U5 OR U3 AVIAN SARCOMA VIRUS LONG TERMINAL REPEATS INFLUENCE INTEGRATION**

The value of reconstituted systems lies in the ability to rapidly analyze mutations that influence integration. In ASV IN, the percentage of nonconcerted integration events can be increased by introducing base changes into the LTR sequences (1, 41, 85), which presumably alter the binding affinity of IN for the LTR recognition sequences. By changing the reaction conditions to favor IN-DNA contacts, the concerted nature of the integration can be rescued and individual integrants can be sequenced (41). When a 4-bp substitution was placed in the ASV U5 LTR, changing CTTCATT to GAAGATT, it resulted in a slight decrease in the efficiency of integration activity compared to that for a donor with a wild-type U5 LTR (1). However, one of every seven integrants sequenced contained deletions in the LTRs. In one case, 10 bases were removed from the U5 LTR, so that IN used the first internal CA dinucleotide for the nucleophilic attack. In a second integrant, IN left the mutation in the U5 LTR but deleted sequences in the wild-type U3 LTR, utilizing the first internal GA dinucleotide to drive the integration reaction (1). This latter result reproduced genetic changes observed in vivo (72) when a mutation placed at one LTR altered the processing of the other. Note also that mutations placed in U3 and U5 have similar effects on integration in terms of specificity. However, mutations placed in U3 have a threefold more deleterious effect on the efficiency of integration in vitro than do the same mutations introduced into the U5 LTR (41, 51, 85).

**RECONSTITUTION OF HUMAN IMMUNODEFICIENCY VIRUS CONCERTED DNA INTEGRATION**

Goodarzi et al. (37) have reported a reconstituted concerted HIV-1 IN-dependent integration system involving a 469-bp donor DNA constructed with the NdeI preprocessed ends. This system still exhibits insertion of two donor DNA molecules into a target, and only about half of the integrants resulted from a concerted DNA mechanism. An HIV-1 mini-donor DNA integration system comparable to the ASV system described by Aiyar et al. (1) has also been developed by using recombinant HIV-1 IN (42). This system appears to approximate integration in vivo more closely. The HIV-1 donor DNA contains only 20 bp of the HIV-1 LTR termini flanking the supF suppressor gene and uses the same target DNA and HMG protein family members as does the ASV system. In contrast to the ASV reconstituted system, where 60% of the sequenced integrants had 6-bp acceptor DNA duplications characteristic of ASV integration in vivo, the duplication of the acceptor DNA at the site of HIV-1 donor integration was almost exclusively 5 bp, characteristic of HIV-1 integration in vivo (42). The differences in base pair duplications in the ASV and HIV-1 reconstituted systems may reflect differences in the stability or conformational characteristics of protein-protein interactions among the respective ASV and HIV-1 IN dimers or tetramers, which form complexes with the ends of the donor and the acceptor DNA. Such differences could influence the spacing of staggered breaks introduced into the acceptor DNA, thereby altering the size of the duplications. The HIV-1 IN-dependent reactions also differ from those of ASV in that less than half or approximately 75% of the HMG-2 and HMG-I(Y) integrants examined, respectively, resulted from a concerted mechanism. The remainder resulted from multiple independent one-ended donor integration events that produce deletions in the target DNA (42). While all of the HMG proteins...
tested stimulated integration in vitro, HMG-I(Y) yielded the most concerted DNA integration products, consistent with the finding of this protein in HIV-1 preintegration complexes (29).

**CELLULAR PROTEINS**

HMG proteins represent a large family of nonhistone DNA binding proteins which are localized primarily in the nucleus of eukaryotic cells and which can be extracted in the presence of acid and salts. The proteins are classified into several families including HMG-1/-2, HMG-14/-17, and HMG-I(Y). The HMG proteins are relatively small, ranging from approximately 11 kDa for HMG-I(Y) and HMG-14/-17 to 25 kDa for HMG-1/-2, and are known to modulate chromatin structure and function (10). HMG proteins have common functional features, including (i) binding to the minor groove of double-stranded DNA; (ii) recognizing DNA structure rather than sequence; (iii) preferentially interacting with bent, supercoiled, or distorted DNA structures; (iv) binding to non-B-form DNA structures such as four-way-junctions and cisplatin adducts; (v) unwinding, bending, and supercoiling DNA substrates in the absence of ATP hydrolysis; and (vi) selectively interacting with other sequence-specific transcription factors as part of gene transcription regulatory complexes.

HMG protein family members increase the efficiency of integration in vitro by acting on the donor DNA (1) without forming stable complexes with IN or the LTR IN recognition sequence, as judged from gel shift and coprecipitation experiments (42). A truncated HMG-I(Y) protein (Δ50–90), which preserves the region of HMG-I(Y) that binds most tightly to substrate DNAs (43), is capable of stimulating integration as well as wild-type HMG-I(Y) does. In contrast, another mutant of HMG-I(Y) (II, III) which has several point mutations preventing protein-DNA interactions while retaining protein-protein interactions, does not stimulate integration in vitro. Taken together, these results imply that HMG-I(Y) needs to associate with the DNA to stimulate integration.

In addition to HMG proteins, another host protein may be essential for in vivo integration. BAF functions as a required factor for efficient integration by preventing auto-integration. Lee and Coffin first noted that MoMuLV was resistant to auto-integration and hypothesized that the lack of auto-integration was due to incomplete reverse transcription (63). Lee and Cragie found that a cellular protein, BAF, prevented auto-integration, and Chen and Engelman demonstrated that BAF could stimulate integration, implying that BAF maintains a competent integration complex by binding the viral DNA into a “open-mesh” complex (15, 61, 62).

**PROSPECTIVE**

With the availability of reconstituted concerted DNA integration assays that closely mimic the genetic complexities of integration in vivo, our understanding of the basic mechanism of integration should dramatically increase. Moreover, these systems will provide the means to screen for drugs targeted at HIV-1 IN, which will potentially open a third avenue of therapy to attack HIV-1-induced AIDS.

**ACKNOWLEDGMENTS**

We thank Mark Andrade, Fox Chase Cancer Center, Philadelphia, Pa., for helpful comments in revising this manuscript and Jerry Alexandratos, Macromolecular Structure Laboratory, NCI-FCRDC, Frederick, Md., for creating the HIV-1 IN structural image.

**REFERENCES**


