Role for a region of helically unstable DNA within the Epstein–Barr virus latent cycle origin of DNA replication oriP in origin function

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Abstract

The minimal replicator of the Epstein–Barr virus (EBV) latent cycle origin of DNA replication oriP is composed of two binding sites for the Epstein–Barr virus nuclear antigen-1 (EBNA-1) and flanking inverted repeats that bind the telomere repeat binding factor TRF2. Although not required for minimal replicator activity, additional binding sites for EBNA-1 and TRF2 and one or more auxiliary elements located to the right of the EBNA-1/TRF2 sites are required for the efficient replication of oriP plasmids. Another region of oriP that is predicted to be destabilized by DNA supercoiling is shown here to be an important functional component of oriP. The ability of DNA fragments of unrelated sequence and possessing supercoiled-induced DNA duplex destabilized (SIDD) structures, but not fragments characterized by helically stable DNA, to substitute for this component of oriP demonstrates a role for the SIDD region in the initiation of oriP-plasmid DNA replication.

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Keywords: Epstein–Barr virus; oriP; DNA unwinding element; EBNA-1; SIDD structures

Introduction

The Epstein–Barr virus (EBV) genome is usually maintained as a single- or multiple-copy plasmid in latently infected cells (Mecsas and Sugden, 1987; Yates, 1996). EBV latent cycle DNA replication may initiate at one of several sites and each plasmid is replicated once per cell division cycle during S phase (Adams, 1987; Little and Schildkraut, 1995; Mecsas and Sugden, 1987; Norio and Schildkraut, 2001; Yates, 1996). The best-characterized EBV latent cycle replication origin is called oriP (Yates et al., 1984). Although this origin is not required for the establishment of the EBV genome as a replicating, extrachromosomal plasmid upon infection of EBV-negative Burkitt’s lymphoma cells (Norio et al., 2000), the presence of oriP within all EBV isolates and the high conservation of its sequence suggests an essential role for this origin in the establishment of EBV latency following the infection of quiescent B lymphocytes in vivo (Koons et al., 2001). A 2.2-kb fragment of EBV DNA encompassing oriP provides cell cycle-regulated replication and stable maintenance of plasmids bearing this fragment (Yates and Guan, 1991) and contains several distinct functional elements (Fig. 1). An element composed of 20 tandemly arrayed binding sites for the EBV nuclear antigen-1 (EBNA-1) designated FR (Family of Repeats) is present at the left end of oriP. FR is required for the maintenance of the EBV genome and plasmids bearing oriP in dividing cells (Harrison et al., 1994; Reisman et al., 1985; Yates et al., 1984) but is dispensable for the initiation of replication within oriP (Harrison et al., 1994; Shirakata and Hirai, 1998; Yates et al., 2000). The replicator of oriP is located approximately 1-kb from FR and is coincident with, or positioned near,
the origin of DNA replication (Gahn and Schildkraut, 1989). The replicator contains four binding sites for EBNA-1, all of which are required for maximal replicator activity (Koons et al., 2001; Yates et al., 2000), and three 9-bp repeats flanking the EBNA-1 sites that are binding sites for the telomere repeat binding factors 1 and 2 (TRF2) (Deng et al., 2002). One or more undefined elements that are not absolutely required for replicator function but increase the efficiency with which replication initiates are located to the right of the EBNA-1 and TRF2 binding sites (Koons et al., 2001).

The binding of EBNA-1 to sites within the replicator bends the DNA at the center of binding but EBNA-1, by itself, is unable to unwind the DNA (Bashaw and Yates, 2001; Frappier and O’Donnell, 1992; Hearing et al., 1992). This information, together with the observations that EBNA-1 is the sole viral protein required for the replication of plasmids bearing oriP (Yates et al., 1985) and that oriP plasmid replication occurs once per cell cycle in synchrony with the duplication of the cellular chromosomes (Adams, 1987; Shirakata et al., 1999), supported the hypothesis that EBNA-1 recruits proteins to the oriP replicator that are involved in the initiation of cellular DNA replication. This prediction was fulfilled in part by the demonstration of the binding of the human origin recognition complex (ORC) and the replication licensing MCM proteins to the replicator of oriP in vivo (Chaudhuri et al., 2001; Schepers et al., 2001). The recruitment of ORC and MCM subunits to oriP was eliminated by deletion of the four EBNA-1 binding sites, the three TRF2 binding sites, and 24 bp of left-hand flanking sequence within the replicator (deletion of EBV nt 8994–9134; (Fig. 1) (Chaudhuri et al., 2001) but the molecular interactions that allow ORC and MCM proteins to associate functionally with the oriP replicator are not known.

In addition to the presence of one or more binding sites for an origin recognition protein that recruits other replication proteins to the origin, the well-characterized origins of the papovavirus SV40 and Saccharomyces cerevisiae, as well as oriC of Escherichia coli, contain elements at, or adjacent to, the origin that facilitate origin unwinding (Kowalski and Eddy, 1989; Lin and Kowalski, 1997; Natali et al., 1992). These DNA unwinding elements (DUEs) are characterized by intrinsic helical instability that may be predicted by a computer program that utilizes the thermodynamic properties of nearest-neighbor dinucleotides (Natali et al., 1992). Some, but not all, DUEs contain sites of hypersensitivity to single-strand specific nucleases in vitro in the absence of the origin recognition protein (Borrowie and Hurwitz, 1988; Kowalski and Eddy, 1989; Lin and Kowalski, 1994; Umek and Kowalski, 1988). These DUEs may be functionally replaced by unrelated sequences that exhibit helical instability (Kowalski and Eddy, 1989; Umek and Kowalski, 1988). We have examined oriP for the presence of potential DUEs using a statistical mechanical method (Benham, 1993) and have identified a supercoiled-induced DNA duplex destabilized (SIDD) structure within oriP. Deletion of this SIDD element greatly reduced the efficiency with which replication initiated within oriP. The replication defect of a plasmid lacking this element was completely rescued by unrelated DNA sequences harboring helically unstable DNA but not by unrelated DNA sequences characterized by high helical stability. These results demonstrate a functional role for a SIDD structure in the initiation of DNA replication within oriP.

**Results and discussion**

Previous analysis of the helical stability of an oriP-bearing plasmid using the thermodynamic properties of nearest-neighbor dinucleotides identified oriP FR as the least helically stable region within oriP (Williams and Kowalski, 1993) and, in the absence of EBNA-1, the single-strand-specific nucleases P1 and T7 cleaved oriP-bearing plasmids within FR (Orlowski and Miller, 1991; Williams and Kowalski, 1993). However, EBNA-1 is bound to the 20 sites within FR throughout the cell cycle and non-base-paired nucleotides have not been detected within FR in vivo or in vitro (Frappier and O’Donnell, 1992; Hearing et al., 1992; Hsieh et al., 1993). To identify potential DUEs within oriP that may not have been identified in previous studies, we analyzed oriP both in the context of the EBV genome and a small recombinant oriP-plasmid (pHEBo-1.1) for the presence of SIDD structures using a statistical mechanical method that measures intrinsic local instability as well as global competition among all sites within a stressed domain and identifies regions within superhelical DNA molecules that are destabilized as well as denatured (Benham, 1992, 1993). The results of this analysis are presented as a destabilization profile in which the incremental free energy G(x) necessary to guarantee denaturation of the base pair at position x is plotted versus sequence location (Fig. 2). High values of G(x) indicate that the bp position is not destabilized by superhelical stresses and, conversely, low values of G(x) identify positions where a base pair is significantly destabilized. The most highly destabilized
region within pHEBo-1.1 is an A + T-rich region within the 3’ region of the β-lactamase gene (around bp position 5800) that has been previously characterized as a site of stable DNA unwinding in supercoiled pBR322 DNA (Kowalski et al., 1988) and the second most highly destabilized region (around bp position 6700) is within the promoter region of the β-lactamase gene (not shown). This latter destabilized region was also shown to be sensitive to a single-strand-specific nuclease in a pBR322 derivative lacking the destabilized region within the 3’ region of the β-lactamase gene (Kowalski et al., 1988). The locations of these regions are consistent with previous findings that sites within duplex DNA predicted to be destabilized by superhelical stresses are associated with cis regulatory loci including gene promoters and terminators (Benham, 1992, 1993). The helix destabilization profile also predicts that a sequence at approximately bp position 1790 (centered at EBV bp position 8660; Fig. 2A) between FR and the replicator is likely to be destabilized by superhelical stresses. This region, as well as oriP, FR, is also predicted to be helically unstable in the context of the EBV genome (Fig. 2B). Neither of these regions within oriP, regardless of sequence context, are predicted to be unwound at a physiological superhelical density of ~0.055. Differences in G(x) for the same region of oriP when in the context of the EBV genome and in pHEBo-1.1 are due to the competition that exists between sites within a single superhelical domain (Benham, 1996).

Early genetic analyses of oriP relied upon the determination of oriP plasmid copy number in stable transformants of EBNA-1-expressing cells as a measure of replication efficiency and concluded that the sequence between FR and the replicator is not absolutely required for replicator activity (Fig. 1) (Reisman et al., 1985). We have found that the measurement of plasmid copy number is not a reliable method for the identification of elements that are not essential for replication to initiate from within oriP but are required for replication to occur with 100% efficiency (i.e., once per cell cycle) (Koons et al., 2001). We therefore analyzed the impact of deletions within oriP upon replication efficiency using a quantitative short-term replication assay to determine if the sequence encompassing the SIDD structure predicted between FR and the EBNA-1/TRF2 binding sites is required for efficient plasmid replication. Test plasmids were co-electroporated into an EBV-positive Burkitt’s lymphoma cell line (Raji) with a plasmid harboring intact oriP and the relative amount of replicated test and control plasmid DNA present within the cells after three cell generations was measured by Southern blotting (Koons et al., 2001). The replication efficiency of an oriP plasmid lacking 642 bp between FR and the EBNA-1/TRF2 binding sites (pHEBo-1.1-Δ8353–8994; Fig. 1) was compared to the replication efficiency of the parental plasmid containing intact oriP (pHEBo-1.1). The positions of relevant restriction enzyme sites within the plasmids used in this study and a representative Southern blot showing the results of this experiment are shown in Figs. 3 and 4, respectively, and the replication efficiencies of the plasmids examined in this experiment are presented in Table 1. Deletion of EBV nt 8353–8994 from pHEBo-1.1 reduced replication efficiency to approximately 30% per cell cycle. The magnitude of the effect of deleting these nucleotides was greater than the effect of deleting an auxiliary element(s) present within EBV nt 9138–9516 from pHEBo-1.1 (Table 1; P < 0.01). Deletion of both nt 8353–8994 and 9138–9516 from pHEBo-1.1 reduced replication efficiency to a level that was indistinguishable from that of a pHEBo-1.1-derivative lacking essential elements of the replicator (pHEBo-2.2; P = 0.33) (Koons et al., 2001).

To address the possibility that helically unstable DNA within EBV nt 8353–8994 contributes to oriP replicator function and its absence is responsible for the replication defect of pHEBo-1.1Δ8353–8994 (Fig. 4 and Table 1), we replaced these nucleotides within pHEBo-1.1 with four fragments of EBV DNA of about the same size. Two of these substitutions were created with DNA fragments of predicted low helical instability (sub1 and sub2; 672 and 650 bp, respectively) and two were constructed with fragments of predicted moderate to high helical instability (sub3 and sub4;
640 and 654 bp, respectively). None of these fragments originated from a region of EBV DNA that contains multiple latent cycle replication initiation sites (Little and Schildkraut, 1995; Norio and Schildkraut, 2001). The destabilization profiles for these DNA fragments within oriP of pHEBo-1.1 are shown in Fig. 5 and the profiles for the sub1 and sub2 DNAs in the context of pHEBo-1.1 and the EBV genome closely resembled one another (Fig. 5 and data not shown). Both sub3 and sub4 were predicted to be more helically unstable when present within the EBV genome (Fig. 5). Plasmids containing sequences with predicted moderate to high helical instability in place of EBV nt 8353–8994 were replicated as efficiently as pHEBo-1.1 (pHEBo-1.1 sub3 and pHEBo-1.1 sub4; Fig. 6 and Table 2). In contrast, plasmids containing sequences with predicted low helical instability in place of EBV nt 8353–8994 exhibited replication defects (compare pHEBo-1.1 sub1 and pHEBo-1.1 sub2 to pHEBo-1.1; P < 0.01 for both comparisons). Similarly, substitution of fragments of predicted high helical instability (sub3 and sub4) for EBV nt 8353–8994 increased the replication efficiency of pHEBo-1.1 Δ8353–8994;9138–9516 to approximately 80–90%, while insertion of fragments of predicted low helical instability (sub1 and sub2) only increased the replication efficiency of pHEBo-1.1 Δ8353–8994;9138–9516 to 30–35% (Fig. 6 and Table 2).

We also replaced the oriP SIDD element centered at EBV nt 8660 with a DNA fragment from the E. coli genome that contains the upstream activating sequence (UAS1) from the ilvPG promoter (Fig. 7) to determine if the well-characterized SIDD element in the ilvPG UAS1 could substitute functionally for the oriP SIDD element (Sheridan et al., 1998; Sheridan et al., 1999). The ilvPG UAS1 was
also predicted to be highly destabilized in pHEBo-1.1/ UAS1 (Fig. 7, panel A) and it replicated as efficiently as pHEBo-1.1 (Fig. 7, panel B).

The ability of a sequence, in which negative superhelicity is predicted to destabilize the DNA by only 2 kcal/mol in several locations (sub4), to substitute functionally for the SIDD region to the left of DS is not surprising when one considers the fact that even relatively small amounts of DNA destabilization can strongly affect the opening process in cases where this process is mediated by a reversible reaction with another molecule. If negative superhelicity destabilizes a region by only 2 kcal/mol (so $G(x) = 8$ kcal/mol), this can drive the equilibrium of the opening reaction more than 25-fold toward the open state, other effects remaining constant. If this opening is the rate-limiting step in a biochemical process, then even small amounts of destabilization can have major effects on the rate of that process. In this regard, it is possible that the modestly destabilized region between EBV nt 8020 and 8353 compensated for the absence of the destabilized region centered on EBV nt 8660 and contributed to the inefficient replication of pHEBo-1.1Δ8353–8994, pHEBo-1.1sub1, and pHEBo-1.1sub2.

The helix destabilization profiles for oriP (Fig. 2) do not predict highly destabilized or denatured sequence elements between EBV nt 9138 and 9516 nor are SIDD structures predicted in this portion of oriP when EBV nt 8353–8994 are deleted from pHEBo-1.1 (data not shown). Therefore, it seems unlikely that the dramatic effect of deleting nt 9138–9516 from pHEBo-1.1Δ8353–8994 upon replication efficiency (Table 1) is the result of deleting another helically unstable region. Instead, these results are reminiscent of the multiple functional elements present within $S. cerevisiae$ chromosomal replication origins (Marahrens and Stillman, 1992) and the origin of DNA replication of SV40 (Parsons et al., 1990), and suggest that functionally distinct elements located on either side of the EBNA-1/TRF2 binding sites within the replicator contribute to the initiation of DNA replication from within oriP.

The ability of four DNA fragments from the EBV genome of similar size but unrelated sequence to increase the replication efficiencies of pHEBo-1.1Δ8353–8994 and pHEBo-1.1Δ8353–8994,9138–9516, albeit to different extents, suggested that the distance between the replicator and an element to the left of the deleted sequence influences replication efficiency. Alternatively, disruption of an EcoRV site to the left of the EBNA-1/TRF2 binding sites in the deletion derivatives of pHEBo-1.1 and its restoration in all four substituted plasmids could have been responsible for the increase in replication efficiency of all the pHEBo-1.1sub1–4 plasmids. These nucleotides are within a sequence (5'-

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**Table 1**

<table>
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<th>Test plasmid</th>
<th>Replication efficienciesa</th>
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<tr>
<td>1</td>
<td>pHEBo-2.2</td>
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<td></td>
<td>pHEBo-1.1</td>
<td>100c</td>
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<td>pHEBo-1.1Δ8353–8994</td>
<td>28 ± 1.6</td>
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<td></td>
<td>pHEBo-1.1Δ9138–9516</td>
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<td></td>
<td>pHEBo-1.1Δ8353–8991</td>
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<td>pHEBo-1.1Δ8353–8991,9138–9516</td>
<td>6.5 ± 1.5</td>
</tr>
</tbody>
</table>

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a Replication efficiencies are the percentage of replicated test plasmid DNA in comparison to replicated reference plasmid (pHEBo-1), normalized to pHEBo-1 (=100%), per cell generation. The values are the mean ± standard deviation and were derived from three independent determinations.

b The replication efficiency of pHEBo-2.2 is reported as <1% as the efficiency was 0.15% in one sample but replicated pHEBo-2.2 DNA was not detected in the other two determinations.

c Replication efficiency over three cell generations relative to pHEBo-1 was 125 ± 10%.

d Replication efficiency over three cell generations relative to pHEBo-1 was 88 ± 11%.
GATAT-3') that is present at a similar position within oriP of rhesus lymphocryptovirus, a highly related gamma-1 herpesvirus (Rivailler et al., 2002). However, restoration of EBV nt 8992–8994 did not improve the replication efficiency of either pHEBo-1.1 derivative lacking nt 8353–8991 (Table 1), suggesting that the spacing between the EBNA-1/TRF2 binding sites and FR or an unidentified element between the right boundary of FR (nt 8029) and nt 8353 is important for oriP function.

Taken together, these results support a role for a region of predicted helically unstable DNA between FR and the replicator of oriP in oriP function. The DNA sequence encompassing this predicted SIDD structure shares several properties with replication origin DUEs. Both are characterized by helical instability and may be functionally replaced by unrelated sequences that also exhibit helical instability (Kowalski and Eddy, 1989; Umek and Kowalski, 1990). Some DUEs, such as the DUEs of S. cerevisiae

**Fig. 5.** Helix destabilization profiles for pHEBo-1.1 derivatives containing foreign DNA substituted for the SIDD element within oriP. The destabilization profiles show the incremental free energy $G(x)$ required to denature the base pair at position $x$ for oriP within pHEBo-1.1 derivatives containing substitutions of helically stable (pHEBo-1.1sub1 [panel A] and pHEBo-1.1sub2 [panel B]) or helically unstable (pHEBo-1.1sub3 [panel C] and pHEBo-1.1sub4 [panel D]) DNA. The positions of FR and DS are indicated by bars over the top of each profile. The approximately 650-bp substitutions are indicated by the dashed lines at the top of each profile. Helix destabilization profiles for EBV nt 133052–133692 (sub3) and 70546–71196 (sub4) in the context of the EBV genome are shown in panels E and F, respectively. The solid bars in panels E and F indicate the DNA fragments used to replace the SIDD element between FR and DS.
chromosomal replication origins and oriC of E. coli are hypersensitive to single-strand-specific nucleases in the absence of binding by the origin recognition protein (Kowalski and Eddy, 1989; Natale et al., 1993), but the unwinding of the DUE of the SV40 origin of DNA replication, called the early palindrome, requires the presence of the origin binding protein (Borowiec and Hurwitz, 1988; Parsons et al., 1990). The sequence around EBV bp position 8660 is not predicted to be unwound (Fig. 2) and is not intrinsically sensitive to single-strand-specific nucleases (Orlowski and Miller, 1991; Williams and Kowalski, 1993). Furthermore, we have been unable to identify in vitro conditions in which the binding of EBNA-1 to oriP induces unwinding of the DNA around EBV bp position 8700. If this sequence serves as a DUE, it is possible that cellular DNA binding proteins are required for its unwinding. Alternatively, this region of oriP may be a component of the replicator that facilitates duplex unwinding at another location. Previous studies of the activation of transcription of the ilvPG promoter of E. coli by integration host factor (IHF) have shown that the binding of IHF within a SIDD structure located upstream of the promoter promotes open complex formation through the transfer of superhelical energy to the promoter (Sheridan et al., 1998, 1999). If the SIDD structure coincides with the physical origin of DNA replication within oriP, it may be concluded from the results.

Table 2

<table>
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<th>Test plasmid</th>
<th>Replication efficienciesa</th>
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<tr>
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<td>pHEBo-1.1</td>
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</tr>
<tr>
<td>pHEBo-1.1sub2</td>
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<tr>
<td>pHEBo-1.1sub3</td>
<td>122 ± 7.1</td>
</tr>
<tr>
<td>pHEBo-1.1sub4</td>
<td>114 ± 6.5</td>
</tr>
<tr>
<td>pHEBo-1.1Δ9138–9516sub1</td>
<td>30 ± 2.1</td>
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<td>pHEBo-1.1Δ9138–9516sub2</td>
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<td>pHEBo-1.1Δ9138–9516sub3</td>
<td>93 ± 7.0</td>
</tr>
<tr>
<td>pHEBo-1.1Δ9138–9516sub4</td>
<td>78 ± 5.2</td>
</tr>
</tbody>
</table>

a Replication efficiencies are the percentage of replicated test plasmid DNA in comparison to replicated reference plasmid (pHEBo-1), normalized to pHEBo-1.1 (= 100%), per cell generation. The values are the mean ± standard deviation and were derived from three independent determinations.
b Replication efficiency over three cell generations relative to pHEBo-1 was 89 ± 11%.

Test plasmid Replication efficienciesa
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<td>pHEBo-1.1Δ9138–9516sub4</td>
<td>78 ± 5.2</td>
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Replication efficiencies are the percentage of replicated test plasmid DNA in comparison to replicated reference plasmid (pHEBo-1), normalized to pHEBo-1.1 (= 100%), per cell generation. The values are the mean ± standard deviation and were derived from three independent determinations. Replication efficiency over three cell generations relative to pHEBo-1 was 89 ± 11%.

Fig. 6. DNA fragments characterized by high helical instability substitute functionally for EBV nt 8353–8994 within oriP. The ability of four fragments of EBV DNA characterized by low (sub1 and sub2) or high (sub3 and sub4) helical instability to substitute functionally for EBV nucleotides 8353–8994 within pHEBo-1.1Δ8353–8994 and pHEBo-1.1Δ8353–8994;9138–9516 was determined as described in the legend for Fig. 3 and Materials and methods, and a representative Southern blot showing the results of one of three determinations is presented in this figure. The test plasmids co-electroporated with pHEBo-1 are indicated above each lane. The three different fragments produced by digestion of EBV DNA with one of three combinations of restriction enzymes used in this experiment to separate the test plasmid from pHEBo-1 and EBV DNA are indicated by arrows on the left side of the figure. The top arrow indicates the position of the EBV genome fragment in lanes 2, 3, 4, 7, and 8. The middle arrow shows the position of the EBV genome fragment in lanes 5, 6, 9, and 10. The bottom arrow indicates the position of the EBV genome fragment in lane 1. The positions of the fragments derived from pHEBo-1 and the test plasmids are indicated on the right side of the figure. The different mobilities of the test plasmid-derived fragments are due to differences in the positions of restriction enzyme sites and not the size of the substituted DNA. The DNA fragments migrating between the fragment derived from pHEBo-1 and the fragments derived from pHEBo-1sub3 (lane 5), pHEBo-1.1sub4 (lane 6), pHEBo-1.1Δ9138–9516sub3 (lane 9), and pHEBo-1.1Δ9138–9516sub4 (lane 10) are pHEBo-1-replicative intermediates.
presented here that other regions within oriP may be used inefficiently in its absence.

Materials and methods

Plasmids

The parental plasmids used in this study, pHEBo-1 and pHEBo-1.1, both contain all of oriP and differ solely in the absence of 342 bp in the portion of the plasmids derived from pBR322 from pHEBo-1.1 that allows the plasmids to be distinguished in short-term replication assays (Hearing et al., 1992; Sugden et al., 1985). pHEBo-1.1Δ9138–9516 and a replication-defective deletion derivative that lacks EBV nt 8995–9137, pHEBo-2.2, have been described previously (Koons et al., 2001). pHEBo-1.1Δ8353–8994 and pHEBo-1.1Δ8353–8994;9138–9516 were derived by deleting 642 bp between the unique MscI and EcoRV sites of pHEBo-1.1 and pHEBo-1.1Δ9138–9516, respectively. Regions of EBV DNA substituted for EBV nt 8353–8994 in pHEBo-1.1 and pHEBo-1.1Δ9138–9516 were obtained by PCR amplification using plasmids harboring restriction enzyme fragments of EBV DNA (B95-8 isolate) as template (Dambaugh et al., 1980). The EBV nucleotide coordinates for these regions are 12,051–12,700 (sub1), 3996–4644 (sub2), 133,052–133,692 (sub3), and 70,546–71,196 (sub4). Together with the addition of nucleotides at the ends of the PCR fragments to facilitate cloning, the inserts in the subplasmids were 672 bp (sub1), 650 bp (sub2), 640 bp (sub3), and 654 bp (sub4). EBV nucleotides 8991–8993 were restored to pHEBo-1.1Δ8353–8994 and pHEBo-1.1Δ8353–8994;9138–9516 by ligating an oligonucleotide linker containing EcoRV ends with the large fragment produced by digestion of pHEBo-1.1 and pHEBo-1.1Δ9138–9516 plasmid DNA with MscI and EcoRV. The ends of an EcoRI–HindIII fragment from pDHWT, containing the E. coli ilvPG UAS1, were repaired with Klenow DNA polymerase and the fragment was ligated with the large EcoRV–MscI fragment of pHEBo-1.1 to create pHEBo-1.1/UAS1.

Short-term replication assays

The replication efficiencies of pHEBo-1.1 derivatives (with the exception of pHEBo-1.1/UAS1) were determined by co-electroporating Raji cells with a reference plasmid that replicates once per cell cycle (pHEBo-1) (Yates and Guan, 1991) and the test plasmid. Electroporated cells were maintained in a constantly dividing state for 72 h at which time low molecular weight DNA was isolated. Samples were digested with DpnI and a combination of other restriction enzymes that produced fragments from the endogenous EBV genome, pHEBo-1 and the amount of probe that hybridized to the plasmid-derived fragments was quantitated with a Molecular Dynamics Storm 860 phosphorimager and ImageQuant software. The relative amount of replicated test and reference plasmid DNA in each sample was corrected for
the ratio of the two plasmids in the sample used for electroporation and the replication efficiencies of the test plasmids were calculated. Statistical analysis of the replication efficiencies was performed by one-way analysis of variance and by using Tukey’s “honestly significant difference” test to make pair-wise comparisons. A detailed description of this assay may be found in Koons et al. (2001).

The replication efficiency of pHEBo-1.1/UAS1 was determined by co-transfecting the EBNA-1-positive cell line 293-EBNA (Invitrogen) with 3 μg each pHEBo-1.1/UAS1 and pHEBo-1.1 using the calcium phosphate co-precipitation method (Graham et al., 1977). A parallel culture was co-transfected with 3 μg each pHEBo-2.2 and pHEBo-1.1. The medium containing the calcium phosphate-DNA co-precipitates was replaced with fresh medium after 16 h at 37 °C and low molecular weight DNA was isolated from the cells 72 h following transfection (Hirt, 1967). DNA isolated from 293-EBNA cells co-transfected with pHEBo-2.2 and pHEBo-1.1 was digested with DpnI, EcoRV, and PstI while DNA isolated from cells co-transfected with pHEBo-1.1/UAS1 and pHEBo-1.1 was digested with DpnI and PstI. The digested samples were analyzed by Southern blotting using a BamHI–MscI fragment bearing oriP FR from pHEBo-1 as the probe as described above.

Analysis of sequences for predicted stress-induced duplex destabilization

Theoretical analysis of the unwinding propensity of the experimental plasmid at various superhelix densities was performed using the technique previously developed by Benham (1992, 1993). This method calculates the statistical mechanical equilibrium distribution of a population of identical, superhelical DNA molecules among all available states of denaturation. It then evaluates the equilibrium probability \( p(x) \) of denaturation at single base pair resolution. It also calculates the destabilization energy \( G(x) \), the incremental free energy required to assure that the base pair at position \( x \) is denatured. Thus, high values of \( G(x) \) occur at positions where the duplex is relatively stable, and low values at easily stress-denatured sites. Positions where \( G(x) \) is near zero are denatured with high probability at equilibrium under the assumed level of superhelicity. The calculation of \( G(x) \) enables one to find fractionally destabilized sites—locations that have a low equilibrium probability of denaturation, but where other processes can drive duplex opening with a relatively small incremental input of energy.

All the free energy and conformational parameters used in these calculations have been experimentally measured, usually in multiple ways and under a variety of environmental conditions. So there are no free (i.e., tunable) parameters in these analyses. Yet their results have been shown to be in quantitative agreement with experiment in all cases where experiments have been performed. They accurately predict both the locations and the extents of destabilization experienced by a DNA molecule of known base sequence, on which a specific level of superhelicity has been imposed.

DNA denaturation behaves differently when it is driven by superhelicity within topological domains than it does when driven by temperature in unconstrained molecules. In the latter case, only near neighbor effects are involved. In the former, the superhelical constraint couples together the behaviors of all base pairs within the domain. This coupling occurs because denaturation of any base pair alters its twist, which changes the partitioning of the superhelicity throughout the domain. In this way, every base pair is affected by the denaturation of any other base pair. This makes the transition behavior of a domain potentially complex and interactive, with the opening of some regions coupled to the reversion back to B-form of others. For this reason, thermodynamic stability is not an adequate predictor of stress-induced destabilization (Benham, 1996).

The calculations reported here were performed using the energy parameters appropriate for the nuclease digestion procedure of Kowalski et al. (1988) at a variety of superhelix densities as described.

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