Multiple homeostatic mechanisms in the control of P1 plasmid replication


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Multiple homeostatic mechanisms in the control of P1 plasmid replication


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Many organisms control initiation of DNA replication by limiting supply or activity of initiator proteins. In plasmids, such as P1, initiators are limited primarily by transcription and dimerization. However, the relevance of initiator limitation to plasmid copy number control has appeared doubtful, because initiator oversupply increases the copy number only marginally. Copy number control instead has been attributed to initiator-mediated plasmid pairing (“handcuffing”), because initiator mutations to handcuffing deficiency elevates the copy number significantly. Here, we present genetic evidence of a role for initiator limitation in plasmid copy number control by showing that autorepression-defective initiator mutants also can elevate the plasmid copy number. We further show, by quantitative modeling, that initiator dimerization is a homeostatic mechanism that dampens active monomer increase when the protein is oversupplied. This finding implies that oversupplied initiator proteins are largely dimeric, partly accounting for their limited ability to increase copy number. A combination of autorepression, dimerization, and handcuffing appears to account fully for control of P1 plasmid copy number.

autorepression | DNA replication control | homeostatic control | plasmid copy number control

From early studies of Escherichia coli and its F plasmid, Jacob et al. (1) proposed that initiation of DNA replication was under positive control of a factor, the “initiator,” that binds to a specific DNA site to set in train the process of replication. Pritchard (2) later argued that positive control alone would not provide dynamic stability and that an “inhibitor” is needed to prevent runaway initiation.

Subsequent studies have shown that inhibition generally works by limiting the activity or availability of initiators. For plasmid ColE1, one of the simplest and best understood replicons, binding of a plasmid-encoded inhibitor RNA to the RNA that primes replication inhibits priming (3, 4). For the E. coli chromosome, several negative regulatory steps prevent the initiator DnaA protein from reinitiating replication prematurely. These include sequestration of DnaA binding sites that are situated at the origin ofreplication and at the dnaA promoter, reduction of free DnaA by binding to new sites created by replication, and accelerated hydrolysis of ATP in the active form of the protein, ATP-DnaA (5). The two initiation factors in Schizosaccharomyces pombe, Cdc18 and Cdc1, are inactivated by phosphorylation and proteolysis at the onset of S phase so that reinitiation of replication cannot take place within the same cell cycle (6).

Although initiator limitation seems widely conserved in replication control, an apparent exception is found in a family of bacterial plasmids controlled by repeated initiator binding sites (iterons). Well studied members of this group include plasmids F, P1, R6K, RK2, pSC101, and pPS0 (7, 8). In these plasmids, saturation of initiator binding to origin iterons allows initiation. Iteron–initiator interactions also underlie negative control of replication.

Many attempts have been made to explain iteron-based control. The first was the initiator–titration model (9). It proposes that after replication, daughter origins compete for the limited amount of initiators, preventing saturation of either origin. However, this proposal overlooked the fact that the number of initiator genes also increases upon replication, leading to an increase in initiator synthesis. Moreover, the initiator genes were found to be transcriptionally autoregulated (10). In plasmid P1, the initiator promoter maps within the origin iterons (Fig. 1A). Thus, initiator binding to the origin also results in promoter repression. Autoregulation normally counteracts protein reduction by titration: as titration reduces the free initiator concentration, autorepression is proportionally relieved and compensates for the reduction (11). To keep titration from counteracting autorepression, it was proposed that the titrated initiators pair with promoter-bound initiators and thereby help maintain the repression (12). This mechanism, now called handcuffing (13), is common among transcriptional repressors that loop DNA, where the titrated repressors, instead of relieving autorepression, actually increase it (14–17).

Handcuffing thus could solve the problem facing initiator titration, but a second observation seemed fatal to the model: Vast increase of initiator supply from constitutive sources increased plasmid copy number by no more than 1.5-fold (18–21). This finding suggested that initiators could not be limiting. Instead it was argued that handcuffing, apart from contributing to autorepression, also causes steric hindrance to replication to prevent reinitiation (13, 18). This view was reinforced when the initiator mutants that increase plasmid number (copy-up mutants) were found to be handcuffing defective (22–25).

Some results on plasmid P1, however, still seem better explained by limited initiator supply. When initiators are supplied in trans at a level only 2-fold below the physiological level, the plasmid copy number declines drastically (10). The copy number also falls when extra iterons are supplied in trans, but this drop can be overcome by supplying extra initiators (18, 26–28). Iteron-mediated control thus can be sensitive to initiator concentration.

Here, we show that selection of copy-up mutants of the P1 initiator RepA, isolated under physiological conditions, yields RepA mutants defective in autorepression but only slightly altered in other properties, including the capacity for handcuffing. Limiting RepA by autorepression thus contributes to copy number control under physiological conditions. We also present a quantitative model to show that initiator dimerization, which autoinactivates the protein (29), is functionally equivalent to autorepression in that they both dampen increase of (active) initiator monomer, which results from increase of copy number. This means of dampening RepA monomer increases works just as well when the protein is supplied from constitutive sources. Thus, the experiments that oversupplied the initiator actually supplied more dimers than monomers, which explains in part the marginal increase in copy number seen previously (18–21).

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results also indicate that restraining plasmid overreplication demands reduction of initiator concentration below that which autorepression or autoinactivation can achieve. Therefore, initiator limitation by autorepression and dimerization is insufficient for replication control, making the additional requirement for a mechanism like handcuffing obligatory. It is the combination of these mechanisms that makes the control most efficient, where one can benefit from the presence of others.

Materials and Methods

**MiniP1 Copy Number.** Plasmid-carrying DH5αlac cells were grown from a single colony to OD₆₀₀ = 0.4, and copy numbers were measured as described in ref. 30.

**RepA Binding in Vivo.** Repression of the repA promoter by RepA supplied in trans was used as a measure of RepA–iteron interactions in vivo (12). The assay system was as described except that the RepA source plasmid, pMVG02, was deleted for the repA gene in a miniP1 plasmid, pSP102 (Fig. 1A) (34). E. coli transformants with mutagenized plasmids were then screened on L agar medium containing 400 μg/ml chloramphenicol, on which cells carrying the WT plasmid formed tiny colonies. Approximately 2% of the transformed colonies were distinctly larger and potentially carried copy-up mutant plasmids. Plasmids were isolated from 17 such colonies, and their repA gene was sequenced. In 12 of 17 cases, single base substitutions were found (Fig. 1B). The remaining five had more than one substitution and were not studied further. All 12 plasmids were functional units of the WT gene in a miniP1 plasmid, pSP102 (Fig. 1A) (34).

**Cross-Linking of RepA.** WT and mutant RepAs were diluted to 10 ng/μl by using RepA dilution buffer [50 mM Tris, pH 7.5/250 mM NaCl/1 mM EDTA/10% glycerol (vol/vol)] and incubated overnight at 4°C. An equal volume of cross-linking buffer (10 mM K-phosphate, pH 7.8/150 mM NaCl/1 mM MgCl₂) was added. To 20 μl of the above mixture, 2 μl of a variously diluted sulfo-bis[2-(succinimido-oxyacarbonyl)ethyl]-sulfone (Pierce) solution in cross-linking buffer was added. The reactions were incubated at room temperature for 10 min, stopped by adding 3 μl of 1 M Tris (pH 7.5), and further incubated for 15 min. Subsequently, 13 μl of 3× SDS sample buffer (pH 6.8) was added. The solution was boiled, chilled, and centrifuged, and the supernatant was run on a 12% Tris-glycine SDS gel. Bands representing RepA were detected by immunoblotting.

**Results**

**Copy-Up Mutants of RepA.** The P1 plasmid replication initiator RepA participates in three specific interactions potentially relevant to the negative control of copy number: autorepression, dimerization, and handcuffing. The autorepression is rather efficient because the repA promoter is repressed ~100-fold when RepA is supplied in trans from a WT P1 plasmid (10). Dimerization is also efficient because purified RepA is predominantly dimeric, although the monomers are required for replication and autorepression (29, 31). Monomerization requires remodeling primarily by chaperones DnaJ and DnaK (32, 33). Only the remodeled monomers bind iterons efficiently, allowing replication and autorepression. The dimers may participate in handcuffing because it happens more efficiently in the absence of chaperones in vitro (31).

To assess the importance of these initiator interactions on plasmid copy number control, new RepA copy-up mutants were isolated without biasing the selection for a RepA phenotype and with RepA expressed from its natural promoter. The repA gene was amplified by using error-prone PCR, and inserted in place of the WT gene in a miniP1 plasmid, pSP102 (Fig. 1A) (34). E. coli transformants with mutagenized plasmids were then screened on L agar medium containing 400 μg/ml chloramphenicol, on which cells carrying the WT plasmid formed tiny colonies. Approximately 2% of the transformed colonies were distinctly larger and potentially carried copy-up mutant plasmids. Plasmids were isolated from 17 such colonies, and their repA gene was sequenced. In 12 of 17 cases, single base substitutions were found (Fig. 1B). The remaining five had more than one substitution and were not studied further. All 12 plasmids showed a moderate increase in copy number (1.2- to 2.1-fold; see Table 1, column 2 from the left). Despite the modest extent of the copy number increase, in contrast to the WT plasmid, all of the

**Surface Plasmon Resonance.** Kinetic constants of RepA–iteron interactions were determined by using the Biacore 2000 instrument (Biacore, Piscataway, NJ). A 31-bp oligonucleotide containing the consensus iteron or nonspecific sequences was biotinylated at one end and conjugated to streptavidin-coated Biacore Sensor Chip SA. The binding buffer contained 50 mM Tris (pH 7.5), 50 mM NaCl, 50 mM KCl, 10 mM MgCl₂, 0.1 mM EDTA, 1 mM DTT, 5 μg/ml poly(dI-dC), and 0.05% (vol/vol) Biacore Surfactant P20. RepA was diluted to 30 nM in the binding buffer and flowed through all of the cells. The reference cell was identical to experimental cells except that it contained the nonspecific oligonucleotide. RepA binding to the reference cell was subtracted to account for nonspecific binding. Data analysis was performed by using the BIAEVALUATION 3.0 software.

**RepA Purification, EMSA, and Ligation Kinetics.** The wild-type (WT) and mutant RepAs (all untagged) were purified identically from an overproducer strain as described for the WT RepA (31). Ref. 31 also describes details of EMSA and ligation kinetics.
the mutant plasmids could transform a bacterial strain containing extra iterons [provided by a miniF clone of PlincA, pALA323 (34)]. In the presence of pALA323, the copy number of the mutant plasmids was 0.3- to 2.2-fold that of the WT (Table 1, column 4). The mutant RepAs thus were considered to be defective in iteron-mediated negative control of replication.

The mutations were located in the same region of repA where previously isolated mutations with the copy-up phenotype had been mapped (Fig. 1B) (30, 35). The previous mutants were selected for proficiency in iteron-binding in the absence of the chaperone DnaJ, but they simultaneously gained the copy-up (and incA insensitive) phenotype. Because of the strong correlation of chaperone-independent DNA binding with copy-up phenotypes, the new mutants were tested for chaperone independence. DNA binding in vivo was tested by RepA’s ability to repress its own promoter when the protein was supplied from a constitutive promoter in trans at concentrations close to those in a P1 lysogen (called 1X). The repression values were comparable with and without DnaJ at least for the first three mutants, similar to those characterized previously (Table 1, columns 5 and 6) (30). These mutants were judged chaperone-independent for iteron binding. Results were similar in ΔdnaJ ΔdnaK (30) and ΔdnaJ ΔcbpA cells [data not shown; CbpA is a functional analog of DnaJ (36)]. Thus, chaperone-mediated monomerization appears to be a rate-limiting step for replication.

Interestingly, the three mutants shown in bold in Table 1 (F120L, K123G and K143E; hereafter called 120, 123 and 143, respectively) were not only chaperone-dependent but also DNA binding defective. When the positions of the mutant amino acids were mapped into a 3D model of RepA–iteron complex, mutants 120 and 123 were found to be in one of the helix–turn–helix regions implicated in DNA binding (Fig. 1C). Mutant 143 was just outside this DNA binding domain. These results are consistent with the DNA binding deficiencies of the mutants. The copy-up phenotype of RepA proteins despite the DNA-binding defect encouraged us to study these three mutants further.

When the three mutant proteins were supplied at 1X in trans, they could not support replication of a miniP1 plasmid deleted for its own repA gene. This result is to be expected because reducing the level of WT RepA to ~0.5X abolishes miniP1 replication (10). When the mutant proteins were produced in cis from their natural promoter, they supported miniP1 replication, probably because of initiator overexpression due to the autorepression defect (Table 1, columns 7 and 8). The level of overexpression depended on the degree of binding deficiency, as would be expected from an autoregulated source. However, the compensation for the binding defect by overexpression is expected to let the copy number approach the WT level, but not to exceed it or to make the replication incA insensitive. The three mutants therefore were examined for other functions in which the initiators are known to participate.

**Table 1. Properties of copy-up RepA mutants**

<table>
<thead>
<tr>
<th>RepA</th>
<th>miniP1/ΔincA copy number*</th>
<th>PrepA repression, † (% in trans)</th>
<th>[RepA] in cis (Western)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>WT/miniF</td>
<td>ΔdnaJ/miniF</td>
<td>WT/miniF + incA</td>
</tr>
<tr>
<td>WT</td>
<td>1</td>
<td>0.7</td>
<td>0</td>
</tr>
<tr>
<td>D55G</td>
<td>1.9</td>
<td>3.0</td>
<td>2.2</td>
</tr>
<tr>
<td>A56T</td>
<td>2.1</td>
<td>1.8</td>
<td>1.5</td>
</tr>
<tr>
<td>A117G</td>
<td>1.3</td>
<td>1.3</td>
<td>0.7</td>
</tr>
<tr>
<td>F120L</td>
<td>1.2</td>
<td>1.1</td>
<td>0.3</td>
</tr>
<tr>
<td>K123G</td>
<td>1.3</td>
<td>1.0</td>
<td>1.2</td>
</tr>
<tr>
<td>K143E</td>
<td>1.5</td>
<td>1.6</td>
<td>1.1</td>
</tr>
<tr>
<td>D152V</td>
<td>1.2</td>
<td>1.3</td>
<td>0.4</td>
</tr>
<tr>
<td>T157S</td>
<td>1.6</td>
<td>3.0</td>
<td>1.3</td>
</tr>
<tr>
<td>D172E</td>
<td>1.2</td>
<td>1.3</td>
<td>0.4</td>
</tr>
<tr>
<td>D180G</td>
<td>1.5</td>
<td>1.2</td>
<td>0.8</td>
</tr>
<tr>
<td>K181E</td>
<td>1.3</td>
<td>1.1</td>
<td>0.4</td>
</tr>
</tbody>
</table>

The three mutants shown in bold were not only chaperone-dependent but also DNA-binding defective. WT also is shown in bold.

*Copy numbers are relative to the WT plasmid pSP102 and are mean values from three cultures started from independent colonies. The average one SD was ~20% of the mean.

†The repA promoter was fused to lacZ and was present in one copy in the chromosome. RepA was supplied in trans from constitutive promoter roughly at physiological concentration. The average one SD was ~12% of the mean.

**Phenotypes of Autorepression-Defective Initiator Mutants in ΔdnaJ Cells.** In the absence of DnaJ, the initiator proteins were overproduced even more than in WT cells, confirming that DNA binding of the mutant proteins is chaperone-dependent (Table 1, columns 7 and 8). The overexpression apparently allowed mutant plasmids to replicate in ΔdnaJ cells efficiently (Table 1, column 3). However, copy number measurements indicated that the WT and mutant RepAs are qualitatively different. In the promoter repression assay, the WT RepA showed only modest defect in ΔdnaJ cells: repression reduced from 97% to 76% (Table 1, columns 5 and 6). Although the protein level increased 3.9× from an autoregulatory source in ΔdnaJ cells, it was apparently not high enough for optimal replication because plasmid copy number fell from 1 to 0.7 (Table 1, columns 2 and 3). The fact that the copy number of the mutant miniP1 plasmids was relatively unchanged in ΔdnaJ cells (Table 1, columns 2 and 3) suggests that there could be weakening of iteron binding specific to autorepression, resulting in increased initiator synthesis. Preliminary evidence to this effect has been obtained (see Fig. 5, which is published as supporting information on the PNAS website). Alternatively, the mutants could be defective in mechanisms that facilitate autorepression. As discussed earlier, handcuffing could be one such mechanism. The mutant proteins therefore were tested for handcuffing.

**Handcuffing Efficiency of RepA Mutants in Vivo.** We used three different assays to measure handcuffing. First, we compared copy numbers of isogenic plasmid monomers and dimers. Because pairing in cis is expected to be more efficient than in trans, the origin copy number in dimer-carrying cells should be lower than in cells carrying monomers. We have shown previously that dimer plasmid copy number is >2-fold lower than that...
of the monomer (25). This finding was confirmed in the present study (Fig. 2A). However, within experimental error, the dimer/monomer plasmid copy number ratio was not significantly different when the miniP1 plasmid (pSP102) carried the mutant genes, indicating that the mutants are handcuffing proficient.

The next assay for handcuffing is based on the principle that pairing of sites in a homodimeric plasmid can separate the DNA into two topologically closed domains, preventing diffusion of transcription-generated superhelical tension from one domain to the other (25). A pBR322 derivative carrying five tandem consensus iterons was used in this assay, and a constant level of RepA (40 nM) was supplied in trans in all of the cases. By this assay, considering the experimental error, only mutant 123 was judged defective in iteron pairing (Fig. 2B). In a previous study, the ratio of positive to negative supercoils decreased ~2-fold when properly oriented and phased with respect to the origin iterons, apparently due to cis-handcuffing between the extra iteron and the origin (31). The extra iteron was ineffective in reducing copy number when previously characterized handcuffing-defective mutants were used (31). In the present study, the WT and the three RepA mutants reduced the plasmid copy number similarly, and, therefore, they were judged handcuffing proficient (Fig. 2C). Taken together, the results indicate that handcuffing is unlikely to be the primary defect of the mutants studied here. Further characterization of the mutants was carried out in vitro where it was easier to adjust for binding differences.

RepA Binding to Origin Iterons in Vitro. We first studied binding in vitro by using DNA fragments containing a single iteron. By using an excess of fragments and the DnaJ and DnaK chaperones, we determined the active fraction of the RepA proteins (37). Protein concentrations were adjusted to compensate for variations in the active fraction. Iteron binding of the mutants was significantly improved when both DnaJ and DnaK were present, but the improvement was less than for WT RepA (Fig. 3A). The results confirmed that the mutants are chaperone-dependent and DNA binding-defective. The KD of binding increased ~2- to 3-fold, mostly due to increased dissociation rates (Fig. 3B).

The binding of the mutants to fragments carrying all five origin iterons was studied in the presence of chaperones (Fig. 3C). The pattern of binding for the mutants differed significantly from that of the WT. The mutants showed fewer retarded bands. Their maximum number (five) corresponded to the number of origin iterons, indicating that the mutants are handcuffing proficient. The WT and mutant RepAs were supplied from constitutive sources in different amounts to achieve similar levels of iteron binding. Copy number of miniP1 without any extra iteron in the presence of 1 X WT RepA was taken as 8.0 (30).

have shown previously that an extra iteron reduces miniP1 copy number by ~2-fold when properly oriented and phased with respect to the origin iterons, apparently due to cis-handcuffing between the extra iteron and the origin (31). The extra iteron was ineffective in reducing copy number when previously characterized handcuffing-defective mutants were used (31). In the present study, the WT and the three RepA mutants reduced the plasmid copy number similarly, and, therefore, they were judged handcuffing proficient (Fig. 2C). Taken together, the results indicate that handcuffing is unlikely to be the primary defect of the mutants studied here. Further characterization of the mutants was carried out in vitro where it was easier to adjust for binding differences.

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The binding of the mutants to fragments carrying all five origin iterons was studied in the presence of chaperones (Fig. 3C). The pattern of binding for the mutants differed significantly from that of the WT. The mutants showed fewer retarded bands. Their maximum number (five) corresponded to the number of iterons present in the probe DNA. Moreover, saturation of binding was difficult to achieve in all three cases, particularly for the mutant 123. Increasing the 123 mutant protein concentration
tants were characterized for their strength of dimerization (31).

Handcuffing of Origin Iterons in Vitro. It is known that the kinetics of intermolecular DNA ligation is enhanced if the DNA molecules are brought into close proximity by interactions between their bound proteins, as in handcuffing (13, 38). Starting with short DNA fragments (a few hundred base pairs), a ladder of ligated multimers was generated (Fig. 4A). When the WT and the mutant RepAs were used at low protein concentration, they caused nearly identical ladder formation, indicating similarity in their handcuffing proficiency (Fig. 4B). At higher protein concentration, the ladder formation was more extensive, except for mutant 123. Increasing protein concentration from 3 to 15 nM did not significantly increase the extent of ladder formation, as was also the case for DNA binding (Fig. 4C). Because only minor differences were observed in the ligation assay, the mutants were considered largely handcuffing proficient.

Dimerization of RepA Mutants in Vitro. Because we have suggested that RepA dimers could be required for handcuffing, the mutants were characterized for their strength of dimerization (31).

From the handcuffing proficiency of the mutants, our expectation was that they would be dimerization proficient. The proteins were diluted to the same final concentration and allowed to equilibrate overnight before exposing to a cross-linking agent (Fig. 4C). No significant differences in dimerization between the WT and the mutants were detected.

In summary, it appears that weakening of iteron binding such that autorepression is specifically compromised (Fig. 5) remains the best explanation for the mutant phenotypes. The present mutants are different from previously characterized copy-up mutants where apparent $K_D$ for iteron binding actually decreases (24).

Discussion Transcriptional autorepression of the initiator gene and initiator inactivation by dimerization are the two well conserved features of iteron-based plasmid replication control systems. They strongly reduce initiator availability, but how they bear on plasmid copy number control has remained unclear. Here, we found that selection of copy-up initiator mutants of plasmid P1 under physiological conditions showed some to be autorepression defective. The mutant initiators were overproduced, which we believe caused the copy number increase. Why, then, does overproduction of WT initiators not cause a proportional increase of plasmid copy number? We argue that initiator inactivation by dimerization is one reason. By using kinetic theory, we show that dimerization dampens the increase of monomers when initiators are overproduced (see Fig. 6, which is published as supporting information on the PNAS web site). The theory also predicts that initiator limitation by autorepression and dimerization, although helpful in dampening plasmid overreplication, cannot be adequate for plasmid copy number control. Additionally, direct control by dimers or handcuffing is required. In fact, the copy number control is best explained if the mechanisms cooperate, as we discuss below.

The goal of replication control is to maintain plasmid copy number within narrow limits. For this process, the replication control mechanisms must be dynamic, meaning that they must respond to an increase in copy number by a decrease in the replication rate. Currently three mechanisms seem to contribute to dynamic responses as follows.

(i) Homeostatic handcuffing: An increase in plasmid copy number increases the plasmid–plasmid handcuffing probability, which sterically inhibits replication.

(ii) Homeostatic monomer–dimer competition: Initiator monomers and dimers promote and inhibit replication, respectively. An increase in plasmid concentration leads to an increase in total initiator concentration, which increases the ratio between dimers and monomers and thereby promotes inhibition over initiation.

(iii) Homeostatic initiator limitation: Autoregulatory mechanisms dampen the increase of total initiator concentration in response to higher plasmid copy numbers, and iteron-mediated titration of the initiator then reduces the free initiator concentration.

These mechanisms also seem to be interdependent where one contributes to mechanically execute the dynamics of the other, as explained below for the case of plasmid P1.

Homeostatic Handcuffing. Handcuffing can in principle provide dynamic control on its own, even if RepA is always present in saturating concentrations. The role of controls on RepA then could be to ensure that the initiator supply is always adequate but not wastefully high. Handcuffing possibly relies on RepA dimerization to link plasmid copies. Both RepA autoregulation and dimerization thus may be necessary to mechanically execute the homeostatic dynamics inherent to handcuffing.
Homeostatic Monomer–Dimer Competition. The binding of RepA monomer is required for replication initiation (29), and the binding of RepA dimer, although weaker, has the potential to inhibit replication (31, 39, 40). The dimers could also participate in replication inhibition without contacting DNA by serving as a bridge between bound monomers (41). Handcuffing then may stabilize dimer interactions as has been found in other systems (14–17). The role of handcuffing then can be to mechanically stabilize dimer interactions as has been found in other systems (41). Handcuffing then may inhibit replication (31, 39, 40). The dimers could also participate in initiator replenishment. Once again, handcuffing thus could help just a smaller increase. As mentioned, initiator titration could number still produce an increase in the initiator concentration, but the autoregulation can only dampen the increase in available initiator number. If it were higher, each plasmid copy number still produces an increase in the initiator concentration, mimicking feedback control without ever “feeding back.” The experiments that greatly oversupplied initiators failed to oversupply the monomer significantly because they only bypassed autorepression-based RepA autoregulation, not dimerization-based RepA autoregulation (18, 19, 21). However, the highly damped copy number increase seen in these experiments may still seem inconsistent with the limited ability of dimerization to dampen the increase in RepA monomer (Fig. 6). Most likely, the other two homeostatic mechanisms become dominant when RepA is supplied in unphysiological excess. The existence of multiple control mechanisms also explains the modest increase of copy number by the present mutants. Relaxing autorepression by mutation would start to increase the plasmid copy number, but the effect would be partly counteracted by preferential accumulation of dimers that would dampen the final increase in plasmid copy number by the other two mechanisms. Having multiple homeostatic mechanisms thus can make the system insensitive to changes in any one of them.

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