High-resolution mapping of architectural DNA binding protein facilitation of a DNA repression loop in *Escherichia coli*

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Edited by Sankar Adhya, National Institutes of Health, National Cancer Institute, Bethesda, MD, and approved April 29, 2015 (received for review January 8, 2015)

Double-stranded DNA is a locally inflexible polymer that resists bending and twisting over hundreds of base pairs. Despite this, tight DNA bending is biologically important for DNA packaging in eukaryotic chromatin and tight DNA looping is important for gene repression in prokaryotes. We and others have previously shown that sequence nonspecific DNA kinking proteins, such as *Escherichia coli* heat unstable and *Saccharomyces cerevisiae* non-histone chromosomal protein 6A (Nhp6A), facilitate lac repressor (LacI) repression loops in *E. coli*. It has been unknown if this facilitation involves direct protein binding to the tightly bent DNA loop or an indirect effect promoting global negative supercoiling of DNA. Here we adapt two high-resolution in vivo protein-mapping techniques to demonstrate direct binding of the heterologous Nhp6A protein at a LacI repression loop in living *E. coli* cells.

DNA looping | lac | architectural protein | Nhp6A | *E. coli*

The local inflexibility of double-stranded DNA limits its bending and twisting over hundreds of base pairs, lengths relevant to DNA biological functions, and interactions with proteins (1, 2). In vitro cyclization kinetics experiments show that the length of DNA most likely to form a circle is ~450 bp, with the probability of smaller circles dropping exponentially with length, as predicted by the worm-like chain polymer model (1). The bending and twisting persistence lengths of DNA (distances over which an initial trajectory is lost because of thermal energy) are both on the order of 150 bp (1, 2).

Although DNA is locally stiff, worm-like chain theory predicts that millimeter-length bacterial genomic DNA spontaneously collapse to coils with volumes of a few hundred micrometers cubed. However, DNA packaging into nucleoids, nuclei, and viruses requires at least 400-fold additional compaction by DNA bending and looping beyond what is achieved by thermal energy (3). Eukaryotic nucleosome formation involves wrapping ~150-bp DNA segments almost twice around histone octamer cores, and DNA segments shorter than one persistence length are also bent and twisted into bacterial repression loops, such as those regulating the lac and gal operons (1, 2, 4–6). Components of the lac operon switch can be reassembled to study DNA looping in vivo, where the β-galactosidase (*lacZ*) gene is controlled by simultaneous binding of the tetrameric lac repressor (LacI) to two operator sequences flanking a promoter. It has been shown that the resulting tight DNA loop inhibits promoter recognition by RNA polymerase (1, 4, 7) (Fig. 1 A and B). Thus, understanding the deformation of stiff DNA molecules is important in biology.

Classic (8–13) and more recent (3, 7, 14–20) experiments have manipulated components of the lac operon in vivo to characterize the biophysics of this switch. Changing the relative spacing and DNA affinities of lac operators, and the concentration of LacI allow modeling of the thermodynamic properties of the switch and the elasticities of the polymer components. One of the mysteries resulting from these analyses is the apparent “softness” of DNA in vivo relative to expectations based on in vitro observations (1). Apparent bend-and-twist flexibilities have been estimated to be two- to sevenfold higher in vivo (8, 9, 21). We are interested in understanding the origin of this apparent DNA softening.

A plausible explanation for DNA softening in cells is the presence of abundant sequence-nonspecific “architectural” proteins with the ability to kink DNA, potentially relieving bending strain (Fig. 1C) (22). Architectural proteins include the bacterial histone-like U93 (HU) protein (Fig. 1D) (23, 24) and the eukaryotic high-mobility group B (HMGB) proteins (Fig. 1E) (25–27). Because they bind and kink DNA (28), such proteins reduce the persistence length of DNA in vitro (29–32) and in simulations (33). Architectural DNA bending proteins may facilitate formation of tight repression loops.

Prior studies have explored the role of architectural proteins in the biophysics of bacterial DNA loops at the lac and gal operons. The Adhya laboratory showed that the bacterial HU protein facilitates gal repression by direct binding to kink the looped DNA (34). Such an effect has never been directly shown for loops anchored by LacI. However, we and others have shown that lac repression is substantially weakened in bacteria lacking repressor (LacI) to two operons (8, 9, 21). We are interested in understanding the origin of this apparent DNA softening.

Double-stranded DNA is one of the stiffest polymers in biology, resisting both bending and twisting over hundreds of base pairs. However, tightly bent DNA loops are formed by proteins that turn off (repress) genes in bacteria. It has been shown that “architectural” proteins capable of kinking any DNA molecule without sequence preference facilitate this kind of gene repression. The mechanism of this effect is unknown for DNA loops involving the well-known *Escherichia coli* lac repressor.

Here we adapt high-resolution protein-mapping techniques to show that an architectural protein directly binds tightly looped DNA to facilitate gene repression by the lac repressor.

Significance

Double-stranded DNA is one of the stiffest polymers in biology, resisting both bending and twisting over hundreds of base pairs. However, tightly bent DNA loops are formed by proteins that turn off (repress) genes in bacteria. It has been shown that “architectural” proteins capable of kinking any DNA molecule without sequence preference facilitate this kind of gene repression. The mechanism of this effect is unknown for DNA loops involving the well-known *Escherichia coli* lac repressor. Here we adapt high-resolution protein-mapping techniques to show that an architectural protein directly binds tightly looped DNA to facilitate gene repression by the lac repressor.
Experimental design. (A) lac promoter construct showing cis elements (−35, −10 elements as magenta circles, Shine–Dalgarno element as black triangle). (B) Repression by LacI tetramer (green circles) via a strained DNA loop in cells lacking the E. coli HU architectural protein. (C) Hypothetical facilitation of DNA looping by yeast sequence-nonspecific architectural protein Nhp6A (red triangle). (D) DNA kinking by Anaebaena HU [PDB ID code 1PS1 (54)]. (E) DNA kinking by S. cerevisiae Nhp6A [PDB ID code 1J5N (235)]. Arrows indicate DNA helix axis trajectory. (F) Model of lac promoter (−10, −35, +1 elements in magenta) captured in a replication loop anchored by LacI tetramer (green) simultaneously binding to upstream (cyan) and proximal (blue) operators. Arrow shows direction of transcription. An Nhp6A architectural protein (red) is indicated near the loop to illustrate scale. (G–J) Experimental lac promoter constructs evaluated here.

Fig. 1. Experimental design. (A) lac promoter construct showing cis elements (−35, −10 elements as magenta circles, Shine–Dalgarno element as black triangle). (B) Repression by LacI tetramer (green circles) via a strained DNA loop in cells lacking the E. coli HU architectural protein. (C) Hypothetical facilitation of DNA looping by yeast sequence-nonspecific architectural protein Nhp6A (red triangle). (D) DNA kinking by Anaebaena HU [PDB ID code 1PS1 (54)]. (E) DNA kinking by S. cerevisiae Nhp6A [PDB ID code 1J5N (235)]. Arrows indicate DNA helix axis trajectory. (F) Model of lac promoter (−10, −35, +1 elements in magenta) captured in a replication loop anchored by LacI tetramer (green) simultaneously binding to upstream (cyan) and proximal (blue) operators. Arrow shows direction of transcription. An Nhp6A architectural protein (red) is indicated near the loop to illustrate scale. (G–J) Experimental lac promoter constructs evaluated here.

protein facilitation of DNA looping. One possibility is related to DNA supercoiling. It has been shown that DNA looping can be stabilized by the unrestrained negative supercoiling typical of bacterial cells (36–39). Supercoiling compacts DNA, raising the local concentration of all DNA sites. Furthermore, DNA supercoiling generates plectonemes where the cost of tight DNA looping is paid by superhelical strain (3, 7, 40). We have shown that deletion of genes encoding various nucleoid proteins, including HU, can change the global superhelical density in Escherichia coli (15). Thus, it is possible that architectural proteins act indirectly to stabilize tight DNA loops by promoting processes that increase global supercoiling.

Here we test the hypothesis that architectural proteins facilitate LacI DNA looping by direct binding to the looped DNA. The model is based on the orientation of the HU-deficient E. coli strain by ectopic expression of the S. cerevisiae Nhp6A (red triangle). (D) DNA kinking by Anaebaena HU [PDB ID code 1PS1 (54)]. (E) DNA kinking by S. cerevisiae Nhp6A [PDB ID code 1J5N (235)]. Arrows indicate DNA helix axis trajectory. (F) Model of lac promoter (−10, −35, +1 elements in magenta) captured in a replication loop anchored by LacI tetramer (green) simultaneously binding to upstream (cyan) and proximal (blue) operators. Arrow shows direction of transcription. An Nhp6A architectural protein (red) is indicated near the loop to illustrate scale. (G–J) Experimental lac promoter constructs evaluated here.

sequence-nonspecific nuclease that can be activated by Ca²⁺ ions (Fig. S1A). Importantly, both forms of Nhp6A functionally complement the lac looping defect in ΔHU cells (Fig. S1B). Expression of these Nhp6A proteins allows mapping of Nhp6A binding to four DNA test constructs (Fig. 1 G–J) integrated into the large F′ episome of E. coli. DNA looping is only expected in the O₁₅/₂₅₂ construct (Fig. 1G) (14), where a pair of lac operators is present. Operators are spaced by −78 bp [an integral number of DNA helical turns, given our consistent observation of 11 bp per turn for this region in vivo (3)] to allow formation of an un twisted loop.

High-resolution mapping of Nhp6A binding was achieved by two methods adapted for the current project. A chromatin immunoprecipitation ex onucleosome ligation-mediated PCR (ChIP-exo-LMPCR) method to map protein binding sites at high resolution on a single E. coli promoter was adapted from a published genome-wide eukaryotic protocol (41, 42). The method is outlined in Fig. S2. Briefly, formaldehyde cross-linking and immunoprecipitation of endogenous epitope-tagged protein–DNA complexes is followed by DNA fragmentation and phage λ exonuclease treatment. Cross-linked proteins are detected as obstacles to processive exonuclease digestion, leaving DNA termini adjacent to the complexes. After cross-link reversal, extension of a gene-specific primer, ligation-mediated PCR, and Southern blotting, detection of the immunoprecipitated protein binding sites is achieved at base pair resolution in sequencing gels.

The chromatin endogenous cleavage LMPCR (ChEC-LMPCR) method was adapted for E. coli analysis by modifying protocols also previously implemented in eukaryotes (43, 44). The method is outlined in Fig. S2. Briefly, formaldehyde cross-linking of an endogenously expressed DNA binding protein fused to MNase is followed by transient Ca²⁺ activation of the nuclease to induce site-specific affinity cleavage of DNA at the site of the bound protein. After reversal of cross-links, capping of nonspecific nicks, polishing of DNA termini, extension of a gene-specific primer, ligation-mediated PCR, and Southern blotting, detection of the MNase fusion protein binding sites is achieved at base pair resolution.

ChIP-exo-LMPCR Mapping of Nhp6A at a Lac Repression Loop. ChIP-exo-LMPCR mapping was applied to four lac constructs (Fig. 1 G–J) to map binding sites of endogenous LacI, the σ₇₀ subunit of E. coli RNA polymerase, and heterologous Nhp6A tagged with a Myc epitope. Results were obtained in the presence and absence of the lac inducer isopropyl-β-D-thiogalactopyranoside (IPTG) and are shown in Fig. 2. Banding patterns in the Southern blot of a representative sequencing gel can be interpreted relative to the flanking diagrams indicating positions of operators (when present, dotted lines in Fig. 2), the −10 and −35 promoter elements, and transcription start point (broken arrow). Maxam–Gilbert chemical DNA sequencing lanes (G, G+A) were used for reference. We first mapped LacI and the σ₇₀ subunit of E. coli RNA polymerase as positive controls before applying the technique to map Nhp6A.

In the absence of specific immunoprecipitation, no signal is seen (Fig. 2, lanes 1–4). In contrast, immunoprecipitation of cross-linked LacI protein followed by exonuclease treatment led to strong banding patterns just upstream of occupied operators in the absence (Fig. 2, lane 5), but not in the presence (Fig. 2, lane 6), of IPTG. Interestingly, the position of exonuclease termination is consistently upstream of LacI bound to the strong O₁₅ operator (Fig. 2, lanes 5 and 9), but largely within the binding site of LacI bound to the weaker O₂ (Fig. 2, lane 5). This finding suggests that the technique detects subtle differences in protein affinity and DNA sequence-dependent cross-linking with formaldehyde. For cases with one or zero operators (Fig. 2, lanes 7–12), LacI binding is weaker, as expected in the absence of cooperative interactions, and there are no distinct exonuclease terminations in the absence of lac operators (Fig. 2, lanes 11–12). Exonuclease termination signals were not observed further upstream or downstream from the lac
promoter. These results for LacI confirmed the sensitivity and specificity of the method.

Results for ChIP-exo-LMPCR mapping of the σ70 subunit of E. coli RNA polymerase are shown in Fig. 2, lanes 13–20. The results confirm expectations: the exonuclease termination signals in the promoter are upstream of the −10 and −35 boxes and strongly IPTG-dependent for the tightly controlled promoter within a repression loop (Fig. 2, lanes 13–14), but less IPTG-dependent when the weak proximal O2 lacks an auxiliary operator (Fig. 2, lanes 15–16), more IPTG-dependent again when the stronger proximal Osym is present (Fig. 2, lanes 17–18), and essentially constitutive in the absence of operators (Fig. 2, lanes 19–20). The IPTG-dependent complementarity between promoter occupation signals because of LacI (Fig. 2, lanes 5–12) vs. σ70 (Fig. 2, lanes 13–20), together with the satisfying position-specificity of the signals, provide unprecedented insight into in vivo protein binding by these factors. These observations demonstrate that ChIP-exo-LMPCR is an effective tool for mapping protein binding. The method was therefore applied to map the binding of Nhp6A.

In the absence of Myc-tagged Nhp6A protein, only background exonuclease termination signals are detected (Fig. 2, lanes 21–24) in ChIP-exo-LMPCR, regardless of the HU status of the cells. In contrast, Myc-tagged Nhp6A creates a very strong pair of exonuclease termination signals just downstream from the −35 box of the lac promoter (Fig. 2, lane 25). We assign these signals to Nhp6A architectural protein bound within the lac loop at this position. This is, to our knowledge, the first such in vivo finding. Note that the strong exonuclease termination signals near the top of the image (Fig. 2, red box in lane 25) are also seen when Osym is occupied by LacI (Fig. 2, upper red box in lane 5). We therefore assign these signals not to Nhp6A, but to LacI cross-linked simultaneously with Nhp6A on the same DNA molecules, acting as a bystander source of exonuclease terminations when Nhp6A is immunoprecipitated. As expected, a corresponding Nhp6A bystander signal is observed in cells expressing Nhp6A when LacI is immunoprecipitated (Fig. S3). The presence of strong exonuclease termination signals attributed to Nhp6A binding within the repression loop correlates with loss of downstream exonuclease termination signals expected for coimmunoprecipitated LacI bound at O2 (Fig. 2, compare lower red box in lane 5 and dotted red box in lane 25). We interpret this suppression as evidence that a large fraction of the captured DNA molecules were cross-linked to LacI at Osym and Nhp6A within the promoter. This finding would explain why most captured DNAs terminated exonuclease cleavage upstream of O2. The strong exonuclease termination signals assigned to Nhp6A within the LacI loop are much attenuated upon gene induction by IPTG (Fig. 2, lane 26), and in the remaining lanes (Fig. 2, lanes 26–32), consistent with the absence of DNA looping in these cases. Thus, formation of the novel Nhp6A complex is strictly dependent on tightly looped DNA.

**ChEC-LMPCR Mapping of Nhp6A at a LacI Repression Loop.** With the described ChIP-exo-LMPCR data showing evidence that the heterologous Nhp6A protein binds directly in LacI loops, we sought to corroborate this result with an independent in vivo protein mapping method. We therefore adapted a ChEC-LMPCR mapping method (43) and applied it to the same four lac constructs to map DNA binding by a Nhp6A–MNase fusion protein. Results are shown in Fig. 3. For the strongly looped Osym/O2 construct, only background signals are observed in the absence of the Nhp6A–MNase fusion (Fig. 3, lanes 1–9). In contrast, a very strong cleavage signal is detected just downstream of the −35 promoter element with increasing Cd2+ activation of the Nhp6A–MNase fusion (Fig. 3, lanes 10–12). This signal is greatly diminished upon IPTG induction (Fig. 3, lanes 13–15), in repressed promoters without DNA looping (Fig. 3, lanes 16–21), and in the constitutive promoter (Fig. 3, lanes 22–24). Three areas of weaker nuclease activity are seen in these cases (Fig. 3, lanes 13–24), corresponding to A/T sequences in the promoter. To interpret this background reactivity, Nhp6A–MNase reactivity on tightly looped DNA (Fig. 3, lanes 25–16) was compared with the same template in the absence of Nhp6A–MNase, treated instead with four increasing concentrations of exogenous MNase after formaldehyde cross-linking in the presence of Nhp6A. The results (Fig. 3, lanes 27–30) confirm that low nonspecific MNase reactivity at A/T
promoter sequences is responsible for the background cleavage signal, whereas the dominant cleavage signal near the center of the lac loop in Fig. 3, lanes 10–12 and 25–26, is a result of direct binding of Nhp6A-MNase in the loop. Endogenous MNase fusion proteins create detectable DNA cleavage even before addition of Ca$^{2+}$ (Fig. 3, lane 25). This is likely because of intracellular Ca$^{2+}$ concentrations (reportedly near 100 nM) in E. coli (45). These ChEC-LMPCR data confirm the results of ChIP-exo-LMPCR (Fig. 2).

A Tightly Bent lac Promoter Sequence Recruits an Architectural DNA Binding Protein. We wished to determine if Nhp6A is recruited to a specific sequence or to a preferred position within the tight repression loop. We therefore compared Nhp6A binding in stable lac repression loops of 67.5 bp, 78.5 bp, and 89.5 bp. Remarkably, Nhp6A binding is detected at the same sequence in all three loops, both by ChIP-exo-LMPCR and ChEC-LMPCR (Figs. S4 and S5). Because the relative position of this sequence is different in each of the loops, this result shows that architectural protein binding is determined by sequence in these tightly bent DNAs.

Summary and Prospects. We previously showed that loss of the abundant E. coli HU architectural protein disables DNA looping by LacI in vivo (14), and that heterologous eukaryotic HMGB architectural proteins can complement this defect (16). Loss of HU reduces the global unstrained negative superhelical density in E. coli (15), an effect that might indirectly reduce DNA looping by LacI (38). On the other hand, tight DNA looping in the gal operon has been shown to be facilitated by direct HU binding in the DNA loop, and fitting to a thermodynamic model of in vivo E. coli LacI looping data, as well as Monte Carlo simulations of LacI loops, also raise the possibility of loop facilitation by direct architectural protein binding. Our data do not rule out a role for global supercoiling effects. However, using two novel adaptations of high-resolution in vivo methods we provide clear evidence of direct DNA loop binding by Saccharomyces Nhp6A, an architectural protein complementing an HU defect in the test strains. The ChIP-exo-LMPCR and ChEC-LMPCR methods provide base pair-resolution data documenting signals because of λ exonuclease and tethered MNase, respectively. Mapping data for LacI, σ$^{70}$, and Nhp6A are summarized in Fig. 4A. The complementing patterns of LacI and σ$^{70}$ binding under repressed vs. induced conditions provide unprecedented insight into protein occupancy of this series of engineered looped and unlooped E. coli promoters and lac operators. The results are entirely consistent with expectations, and also reveal the potential sensitivity of ChIP-exo-LMPCR to subtle effects of operator sequence and affinity on patterns of formaldehyde cross-linking and exonuclease termination. This sensitivity can complicate mapping of precise protein binding sites (42). The method is also shown to detect cross-linked bystander proteins coinmunoprecipitated with local target proteins of interest.

Of greatest significance to testing the current hypothesis is the clear detection of the Nhp6A protein bound to a specific lac promoter sequence within the DNA looped by LacI (Fig. 4A). Exonuclease termination sites just downstream of the −35 box of the test promoter suggest a single Nhp6A binding site in the loop (Fig. 4B). Comparison of the apparent Nhp6A binding sequence with the reported high-resolution Nhp6A-DNA NMR structure (25) immediately suggests a model for this interaction (Fig. 4B and Fig. S6). Nhp6A binding appears to map to a 5′−TG/CA base pair step in the lac promoter, a sequence known to be readily kinked (46). Nhp6A kinking at such a dinucleotide involves intercalation of methionine 29. We propose that this kinkable site within the lac promoter plays a natural role in recruiting architectural protein stabilizers of the tight repression loop as has been suggested for HU (20). We show using ChIP-exo-LMPCR and ChEC-LMPCR that this is the preferred Nhp6A binding sequence in repression loops of three different sizes, and that a loop lacking this sequence does not recruit Nhp6A. Furthermore, Nhp6A binding is detected only in cases of tight DNA looping by LacI. Thus, we show that the sequence of the lac promoter encodes an architectural DNA binding protein site reminiscent of the architectural protein binding site at the gal loop.

Efforts are underway to extend the present methods to map binding of the endogenous heterodimeric HU protein where recent in vitro, in vivo, and simulation studies propose repression facilitation by direct loop binding (34, 35). In addition, the constructs studied here are based on components of the lac operon control switch, but differ in subtle sequence and spacing details
relative to the wild-type lac promoter. We are now mapping protein binding sites on the wild-type lac operon promoter in vivo.

Our experimental results are directly relevant to the fascinating recent experimental and simulation work of the Phillips and coworkers (20) and Olson and coworkers (35). Boedicker et al. (20) confirm our prior in vivo result (14) that E. coli architectural protein HU facilitates gene repression by LacI, and extend the work in vitro using tethered particle motion experiments. The authors show that DNA looping sequence effects become latent in the presence of HU. Using data fitting to an insightful statistical mechanics model, Boedicker et al. (20) go on to propose that loop facilitation is because of two HU proteins binding directly within the lac loop. The absence of supercoiling effects in the tethered particle motion experiments tends to support this direct loop binding hypothesis. Wei et al. (35) use Monte Carlo simulations to argue that, at equilibrium, sequence-nonspecific architectural proteins, such as HU, will spontaneously decorate tight DNA loops at preferred positions due to thermodynamic effects (minimizing the free energy of the strained system). The authors simulate random uptake of HU proteins onto the 92-bp wild-type lacI loop and find one or two HU proteins bound in cases of successfully closed structures. In light of these predictions and the data presented here, it will be very interesting to apply the ChIP-exo-LMPCR and ChEC-LMPCR methods to experimentally map HU binding sites on the wild-type lacI loop in living bacteria. These experiments promise important new insights into the mystery of apparent DNA softening in vivo.

Methods

Bacterial Strains. The four DNA promoter/operator looping DNA constructs (Fig. 1 G–J) used in this study were based on plasmid pJ992 (14) created by modification of pFW11-null (47). See SI Methods for full details.

Protein Expression Constructs. Nhp6A and Nhp6A-MNase protein expression constructs were created by inserting purified PCR products into plasmid pJ1035, a modified version of pLX20 containing a promoter driving moderate levels of protein expression (14). Both full-length Nhp6A (pJ1327) and Nhp6A Δ2–12 (pJ1328) were previously described (16). See SI Methods for full details.

Molecular Modeling. Molecular docking and graphics were implemented with 3D-DART (48) and Pymol (49).

β-Galactosidase Enzyme Assays. A liquid β-galactosidase colorimetric enzyme assay measured lacZ expression was performed as described previously (15). The repression ratio (RR) is given as the ratio of induced/repressed expression, where induction is obtained by addition of 2 mM IPTG. Analysis of the resulting lac reporter gene expression patterns was performed as described previously (50), with fitting optimization using a simplex and inductive search hybrid algorithm (51).

Bacterial Growth and Formaldehyde Cross-Linking. E. coli strains carrying the indicated protein expression plasmids were grown to log phase in 40 mL LB medium at 37 °C in the presence or absence of 2 mM IPTG. Cultures were pelleted at 4,000 × g for 10 min at room temperature and resuspended in 20 mL PBS (Mg²⁺- and Ca²⁺-free) before cross-linking of macromolecules by the addition of 3% (vol/vol) formaldehyde (Sigma) to a final concentration of 0.75%. Cultures were maintained at room temperature with constant gentle swirling for 20 min. Cross-linking was terminated by addition of cold, 2 M Tris-HCl (pH 8.0) to a final concentration of 260 mM. Cells were harvested by centrifugation, washed three times with 4 mL cold PBS, and cell pellets stored at −80 °C until ready for processing.

ChIP-exo-LMPCR Analysis. This method was adapted for bacterial analysis based on previous eukaryotic methods (41, 42). See SI Methods for full details.

ChEC-LMPCR Analysis. This method was adapted for bacterial analysis from previous publications (43, 44). See SI Methods for full details.

LMPCR. λ Exonuclease and MNase cleavage sites were analyzed by adaptation of standard LMPCR methods (52, 53). See SI Methods and Fig. S7 for full details.

ACKNOWLEDGMENTS. The authors thank Justin Peters for expert assistance. This work was supported by the Mayo Foundation and National Institutes of Health Grant GM75965 (to L.J.M.).
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