Acidic C-terminal tail of the ssDNA-binding protein of bacteriophage T7 and ssDNA compete for the same binding surface

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ssDNA-binding proteins are key components of the machinery that mediates replication, recombination, and repair. Prokaryotic ssDNA-binding proteins share a conserved DNA-binding fold and an acidic C-terminal tail. It has been proposed that in the absence of ssDNA, the C-terminal tail contacts the ssDNA-binding cleft, therefore predicting that the binding of ssDNA and the C-terminal tail is mutually exclusive. Using chemical cross-linking, competition studies, and NMR chemical-shift mapping, we demonstrate that: (i) the C-terminal peptide of the gene 2.5 protein cross-links to the core of the protein only in the absence of ssDNA, (ii) the cross-linked species fails to bind to ssDNA, and (iii) a C-terminal peptide and ssDNA bind to the same overall surface of the protein. We propose that the protection of the DNA-binding cleft by the electrostatic shield of the C-terminal tail observed in prokaryotic ssDNA-binding proteins, ribosomal proteins, and high-mobility group proteins is an evolutionarily conserved mechanism. This mechanism prevents random binding of charged molecules to the nucleic acid-binding pocket and coordinates nucleic acid–protein and protein–protein interactions.

S
ingle-stranded DNA (ssDNA)-binding proteins are a key component of the machinery for replication, recombination, and DNA repair (1). Once assigned to such mundane roles as eliminating secondary structure in ssDNA and protecting DNA from cleavage by nucleases, they are now emerging as key components in coordinating reactions at replication forks (2, 3).

The structures of several prokaryotic ssDNA-binding proteins have been solved (4–9). Although these proteins do not have sequence homology, their structures share a common oligosaccharide/oligonucleotide-binding fold (OB-fold), suggesting that they have a similar mode of action. The OB-fold is comprised of antiparallel β-sheets forming a barrel with a well defined cleft. Structural and mutagenesis data have shown that ssDNA binds within their cleft via stacking and electrostatic interactions. Several invariant aromatic residues surrounded by positively charged amino acids are found in all ssDNA-binding proteins. Although the aromatic residues stack with the DNA bases, the positively charged amino acids contact the phosphate backbone (4–8, 10, 11).

In addition to a common structure, all prokaryotic ssDNA-binding proteins have an acidic C-terminal tail. This C-terminal tail is essential for DNA replication, and phage and bacteria lacking the C-terminal tail fail to grow (11–15). In vitro, ssDNA-binding proteins lacking the C-terminal tail have a higher affinity for ssDNA (12, 15–21). The characterization of the nucleic acid-binding properties of gp32, the ssDNA-binding protein of phage T4 (17–20), led to the proposal of a model for the binding of gp32 to polynucleotides (Fig. 1). The model postulates that the negatively charged C-terminal tail binds electrostatically to the positively charged DNA-binding cleft and that it is displaced upon binding of DNA. Thus, proteins lacking the C-terminal tail have a vacant DNA-binding cleft lacking the electrostatic shield provided by the C-terminal tail, and ssDNA can bind directly to the cleft without having to compete with the C-terminal tail for occupancy of this site.

The finding that the C-terminal tail of prokaryotic ssDNA-binding proteins mediates multiple protein–protein interactions at the replication fork in Escherichia coli and in bacteriophage T7- and T4-infected cells further extended the model presented in Fig. 1 (2, 11, 22–24). It was proposed that the C-terminal tail functions as a two-way switch that coordinates and modulates the ssDNA binding and the protein–protein interactions. When the ssDNA-binding protein is bound to ssDNA, then the C-terminal tail is free to electrostatically interact with other proteins and recruit them at the site of the ssDNA and/or modulate their activity (5).

The gene 2.5 protein (gp2.5) is one of the four proteins that comprise the T7 replisome. The three other proteins are the T7 gene 5 DNA polymerase; its processivity factor, E. coli thioredoxin (trx); and the multifunctional gene 4 helicase-primase. The acidic C-terminal tail of gp2.5 is critical for the interactions of the protein with T7 DNA polymerase and helicase-primase and was evidenced by their abolishment when the tail is deleted (11, 23). Recently, we have shown that the C-terminal tail of gp2.5 binds to a highly positively charged segment located in the thumb subdomain of the T7 gene 5 DNA polymerase (25). This fragment also is the site of binding of the processivity factor, E. coli trx, and the C terminus of gene 4 helicase-primase. Thus, the C-terminal tail of gp2.5 could potentially interfere with all interaction pairs at the T7 replication fork: directly by being part of the contact interface itself or indirectly by competing with another partner for the same contact surface (11, 23, 26, 27). The multiple interactions of the C terminus of gp2.5 could thus function as one mechanism to coordinate the multiple reactions occurring at the replication fork. Accordingly, gp2.5 is critical for establishing coordinated leading and lagging strand DNA synthesis (28–30).

Recently, we showed that, in addition to its acidic charge, a C-terminal aromatic residue on the tail also is essential for the function of gp2.5. gp2.5 lacking its C-terminal phenylalanine is defective in DNA replication and is unable to grow in E. coli (31). In vitro, gp2.5 is defective in its interaction with T7 DNA polymerase in vitro. Interestingly, the T7 helicase-primase also has an acidic C-terminal tail with a functionally important C-terminal aromatic residue (32). The common features of the C-terminal tails of both proteins provide additional evidence that they are part of a regulatory mechanism involving competition for binding to the same site.

gp2.5 is a dimer in solution. It has been proposed that the C-terminal tail of each monomer could bind in the ssDNA-binding cleft of the other subunit in trans, thus stabilizing the dimer interface, seen in the crystal structure of gp2.5 lacking the C-terminal tail (5). The displacement of the C-terminal tail upon

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ssDNA binding was in turn proposed to result in the destabilization of the dimer interface, which could lead to protein monomerization (33). Despite the ample evidence that the C terminus of gp2.5 is critical for its function, there is no information about the structure of the tail or its position with respect to the rest of the gp2.5 molecule. Unfortunately, no such information is available for any other acidic C-terminal tail of prokaryotic ssDNA-binding protein.

In the present study, we use gp2.5 as a model system for a prokaryotic ssDNA-binding protein to test the prediction that the acidic C-terminal tail and ssDNA compete for binding to the ssDNA-binding cleft of the molecule. By employing a combination of chemical cross-linking, competition studies, and NMR chemical shift mapping, we demonstrate that the C-terminal peptide of gp2.5 and ssDNA bind to the same overall surface of gp2.5 in a mutually exclusive manner.

Results

The C-Terminal Tail of gp2.5 Cross-Links to the Core of the Protein. The C-terminal tail of gp2.5 contains multiple amino acids with carboxylic side chains, whereas its predicted site of binding, the DNA-binding cleft, contains multiple residues with amino groups. The 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (EDC)/N-hydroxysulfosuccinimide (sulfo-NHS) cross-linking system (Pierce) activates free carboxyl groups for subsequent reaction with amino group side chains in amino acids (34). Therefore, we chose this cross-linking system because it utilizes the natural chemical composition of the regions of interest of gp2.5 and does not lead to the incorporation of additional components among the cross-linked products.

A peptide having a sequence identical to the C-terminal 26 amino acids of gp2.5 (ASKPRDEESWDEDDDEESEAEDEGDGF) readily cross-links to residues in wild-type gp2.5 (Fig. 2A, lane 2). In these experiments, the cross-linked species are identified by their decreased mobility during electrophoresis through SDS gels. A control peptide containing the same amino acids, but in random arrangement, does not cross-link to residues in gp2.5 (Fig. 2A, lane 3). The same cross-linking reaction was performed with gp2.5Δ26 lacking the C-terminal tail (5). Gp2.5Δ26 behaves similarly to wild type in the cross-linking reaction, producing a cross-linked product with decreased electrophoretic mobility (Fig. 2B). The products of the cross-linking reaction between gp2.5Δ26 and the C-terminal peptide migrate on SDS/PAGE very similarly, if not identically, to full-length gp2.5, demonstrating that only one peptide cross-links to each molecule of gp2.5Δ26. Taken together with the finding that the random peptide does not cross-link to gp2.5Δ26 even when it is in large molar excess (Fig. 2B, lanes 6–8), this observation strongly suggests that the cross-linked product most likely reflects the natural binding position of the acidic C-terminal tail.

The ability of the activated peptide to compete with the native C-terminal tail and cross-link argues that the C-terminal tail is only transiently and weakly bound to the core of the protein. We chose to perform all subsequent experiments by using gp2.5Δ26 to avoid any competition between the natural C-terminal tail and the C-terminal peptide supplied in trans.

ssDNA Abolishes the Cross-Linking of the C-Terminal Peptide to gp2.5. If indeed the C-terminal tail of gp2.5 binds in the ssDNA-binding cleft of the protein, ssDNA should compete with the C-terminal peptide for cross-linking to the protein. gp2.5Δ26 was preincubated with ssDNA (5.4-kb circular ssDNA from bacteriophage ΦX174) for 10 min before activated peptide was added to the reaction. The addition of increasing amounts of ssDNA resulted in abolishment of cross-linking (Fig. 3A). Cross-linking was abolished at 560 μM (nt) ssDNA (Fig. 3A, lane 7).

Cross-Linked Species Do Not Bind ssDNA. The cross-linking experiments shown in Fig. 3A suggest that the C-terminal tail of gp2.5 binds in the ssDNA-binding cleft of the protein. Therefore, the cross-linked species in which the C-terminal peptide is presumably covalently bound in the DNA-binding cleft are expected not to be able to bind ssDNA. To test this hypothesis, we have examined the ability of gp2.5Δ26 cross-linked to peptide to bind to ssDNA by using an ssDNA cellulose column. The cross-linking reaction loaded on the column (Fig. 3B, lane 1) contains gp2.5Δ26 and gp2.5Δ26-peptide cross-linked complex. After two washes with 200 μl of low-salt buffer each, the bound proteins were eluted with 200 μl of high-salt buffer containing 1 M sodium chloride. The binding
proteins by precipitation with trichloroacetic acid (TCA) and resuspended in PAGE-loading buffer, and the products were resolved on 4 –20% SDS/PAGE gels. This experiment exploits this phenomenon to determine whether the same, different, or partially overlapping sets of residues of the 15N-labeled protein are involved in the interactions with different ligands (for a recent review, see ref. 35).

Binding of the C-terminal peptide was in fast exchange on the NMR time scale (peaks gradually moving upon addition of increasing amounts of peptide), indicative of a fast dissociation rate. The binding of the 23-mer ssDNA to gp2.5Δ26 appeared stronger than that of the peptide: in fast-to-intermediate exchange regime on the NMR time scale, with some peaks becoming broadened or completely disappeared (intermediate exchange), indicative of a slower dissociation rate. Upon adding increasing amounts of ssDNA, most of the spectrum of gp2.5Δ26 disappeared (data not shown) presumably due to the formation of high-molecular-weight complexes composed of multiple gp2.5Δ26 molecules bound to the same ssDNA molecule. Therefore, an early titration point with a sub-stoichiometric amount of ssDNA, where the spectral quality was still good, was used for analysis. Examples of peaks with significant chemical shift changes are shown in Fig. 4. Statistical analysis of the observed chemical shifts showed that only a small subset of peaks (Fig. 5, depicted in blue and red) exhibit changes beyond the average background chemical shift change plus two standard deviations. Nine such peaks were observed in the titration with ssDNA (Fig. 5A), whereas 10 such peaks were observed in the titration with gp2.5 C-terminal peptide (Fig. 5B). Seven of those peaks are identical between the two titrations (depicted in blue), showing that the titration of ssDNA and gp2.5 C-terminal peptide affected the same residues from the core module of gp2.5 protein. Because backbone resonance assignments are not available for gp2.5Δ26, NMR could not be used to identify the specific residues of gp2.5Δ26 affected by the interactions. This finding directly demonstrates that ssDNA and the C-terminal tail bind in the same overall area of the surface of the protein. Furthermore, ssDNA binds to gp2.5Δ26 tighter than the C-terminal peptide does, as evidenced by the different magnitude of the chemical shift changes observed upon binding of both ligands (Fig. 5).

Discussion

Organisms from all domains of life and many prokaryotic and eukaryotic viruses encode ssDNA-binding proteins as part of the molecular machinery that maintains the integrity of the genome and ensures its replication. Despite the absence of significant

![Fig. 3. ssDNA-binding and peptide cross-linking are mutually exclusive. (A) Gp2.5Δ26 (4 μM) was preincubated with increasing amounts (56 – 1,120 μM in nucleotides) of 4X174 circular ssDNA for 10 min on ice. EDC/sulfo-NHS-activated peptide (80 μM) was added, and the reaction was incubated at room temperature for 20 min. The cross-linking reactions were stopped with SDS/PAGE-loading buffer, and the products were resolved on 4 –20% SDS/PAGE (Bio-Rad). (B) Wild-type C-terminal peptide (80 μM) was activated with EDC/sulfo-NHS and cross-linked to 4 μM gp2.5Δ26. One half of the cross-linking reaction was loaded on an ssDNA spin column, subjected to two low-salt buffer washes, and eluted with high-salt buffer as described in Materials and Methods. Half of the cross-linking reaction was loaded on gel as a reference for the species subjected to ssDNA cellulose binding (lane 1). The species of gp2.5Δ26 cross-linked to ssDNA cellulose, whereas cross-linking reagents do not bind the column and appear in the low-salt eluate. In contrast, free gp2.5Δ26 protein is able to bind to ssDNA cellulose, whereas cross-linked species do not bind the column and appear in the low-salt wash. This finding further demonstrates that the binding of ssDNA and the C-terminal tail is mutually exclusive.

NMR Chemical Shift Mapping of the Binding Surfaces of the C-Terminal Peptide and ssDNA. We aimed to test whether the C-terminal peptide of gp2.5 and ssDNA bind to the same surface of the protein by performing a series of NMR chemical shift perturbation experiments to monitor the interaction of gp2.5Δ26 with wild-type C-terminal peptide and ssDNA, respectively. In the 1H, 15N heteronuclear single-quantum coherence (HSQC) NMR spectra (Fig. 4), each NH group in the 15N-labeled protein gives rise to a peak in the spectrum at the intersection between the chemical shifts of the respective 15N and 1H nuclei. The exact position of each peak (the chemical shifts of the 15N and 1H nuclei) is influenced by the environment around the NH group in the protein. Any change in this environment (e.g., binding of a ligand in the vicinity of the NH group) affects the peak position. The NMR chemical shift perturbation experiment exploits this phenomenon to determine whether a ligand binds to the labeled protein. If more than one ligand binds to the same protein, NMR chemical shift perturbation experiments can be used to determine whether the same, different, or partially overlapping sets of residues of the 15N-labeled protein are involved in the interactions with different ligands (for a recent review, see ref. 35).

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Discussion

Organisms from all domains of life and many prokaryotic and eukaryotic viruses encode ssDNA-binding proteins as part of the molecular machinery that maintains the integrity of the genome and ensures its replication. Despite the absence of significant
sequence homology, all ssDNA-binding proteins share a common structural core, the OB-fold. In addition, prokaryotic ssDNA-binding proteins, mitochondrial ssDNA-binding proteins, and crenaceous archaeal ssDNA-binding proteins have a flexible acidic C-terminal tail (10).

Earlier studies of the nucleic acid-binding properties of full-length and proteolytic fragments of gp32, the ssDNA-binding protein encoded by bacteriophage T4, led to the proposal that, in the absence of DNA, the acidic C-terminal tail binds in the DNA-binding cleft (17). Studies of E. coli and T7 and T4 replisomes revealed that the C-terminal tails of the cognate ssDNA-binding proteins mediate multiple protein–protein interactions (reviewed in refs. 2 and 3). Thus, the current working model (Fig. 6 A) for the role of the acidic C-terminal tail is that, in the absence of DNA, it binds to the DNA-binding cleft. Upon DNA binding, the tail is displaced and available for protein–protein interactions. Although the model depicts the C-terminal tail of a gene 2.5 monomer binding within its own DNA-binding cleft, it also is possible for the C-terminal tail of one monomer to bind within the DNA-binding cleft of another monomer to form a dimer, as suggested previously (5, 33).

One prediction from the described model is that the binding of ssDNA and the C-terminal tail to the DNA-binding cleft is mutually exclusive. Using cross-linking and affinity binding, we have demonstrated that the cross-linking (i.e., binding of the acidic C-terminal tail to the core of the gp2.5) prevents the protein from binding to ssDNA. Vice versa, the binding of ssDNA competes successfully (i.e., prevents the binding/cross-linking of a peptide), corresponding to the acidic C-terminal tail. Furthermore, NMR chemical shift mapping demonstrates that both ssDNA and C-terminal peptide contact gp2.5Δ26 in the same area, providing additional evidence that indeed the C-terminal tail binds into the DNA-binding cleft of the protein. The range of the changes in the NMR spectra observed upon titration of ssDNA and wild-type peptide demonstrates that ssDNA binds to gp2.5Δ26 with higher affinity than the C-terminal peptide, a result that is consistent with the C-terminal tail being displaced upon ssDNA binding. In summary, using gp2.5 of bacteriophage T7, we demonstrate that the binding of ssDNA and the acidic C-terminal tail to the core of gp2.5 is mutually exclusive as predicted.

The competition of the DNA and the C-terminal tail for the DNA-binding cleft represents an elegant mechanism that ensures efficient coverage of any ssDNA arising in the cell without posing a road block to the DNA processing machinery. All ssDNA-binding proteins studied up to date are abundant in the cell, thus ensuring the timely coverage of any ssDNA arising in the cell. However, the moderate affinity for ssDNA allows prompt removal of the protein upon the recruitment of the DNA processing machinery that would convert the ssDNA to dsDNA via replication, recombination, or repair. The relatively high cellular concentration of ssDNA-binding protein increases the probability that it will bind randomly to any negatively charged molecule or surface in the cell. Prokaryotic ssDNA-binding proteins avoid that possibility by using their acidic C-terminal tail as a protective electrostatic shield. We propose that the C-terminal tail oscillates between the bound and the unbound position, thus sensing the presence of ssDNA in the cell (Fig. 6 A).

The oscillation mechanism also can explain why the C-terminal tail can mediate the interaction of gp2.5 with T7 DNA polymerase and helicase-primase in the absence of DNA (11, 23). In the presence of ssDNA, the C-terminal tail is displaced from the DNA-binding cleft because of the higher affinity of the protein for ssDNA. It is possible that the binding of the tail is purely electrostatic, whereas the DNA binding employs electrostatic interactions with the backbone and stacking interactions with the bases, collectively resulting in a stronger interaction.

Further support for the electrostatic shield hypothesis comes from the finding that T4 gp32 can bind dsDNA and RNA, although with much lower affinity (i.e., when negatively charged molecules are available in sufficient concentration, they bind to the positively charged cleft) (16–18, 20, 36, 37). Similar to ssDNA, the removal of the acidic C-terminal tail results in higher affinities for these substrates. Deletion analysis of the C-terminal tail of gp2.5 demonstrated that the gradual removal of charged residues results in the gradual increase of the affinity for ssDNA (31). This observation provides further evidence that the C-terminal tail shields the ssDNA-binding cleft.

Interestingly, T4 gp32 binds specifically to a translational operator in its own mRNA and self-regulates its expression, thus allowing effective control of the abundance of the protein (38). No data are available about such control mechanism for any other ssDNA-binding protein. However, it is logical to expect that the abundance of these proteins should be regulated to ensure the availability of sufficient amounts of protein to cover any transiently arising ssDNA without posing a road block to the DNA processing machinery. However, the electrostatic shield of the acidic C-terminal tail provides a means of protection of the DNA-binding cleft from binding to random charged surfaces.

Similar to prokaryotes, eukaryotes also employ OB-folds as modules of their ssDNA-binding proteins, but their molecules lack an acidic tail. The best studied eukaryotic ssDNA-binding protein, human replication protein A, is composed of several subunits, harboring one or more OB-fold modules (39). Most likely, the heteromeric complex of RPA has a sophisticated 3D structure that prevents binding of random charged molecules to the DNA-binding cleft(s) of the protein complexes. Also, the presence of multiple OB-folds within the same complex creates multiple possibilities for modulation of the DNA-binding activity.

The presence of phage and bacterial ssDNA-binding proteins together in the cell during infection is an interesting biological...
Fig. 6. Model for the functional role of the acidic C-terminal tail. (A) The C-terminal tail functions as an electrostatic shield. In the absence of ssDNA, the C-terminal tail oscillates between the bound and unbound states, thus effectively protecting the positively charged DNA-binding cleft from binding to random negatively charged surfaces and sensing for the presence of ssDNA. For clarity, the figure represents only one monomer of a dimer of gp2.5 with C-terminal tail binding within its own DNA-binding cleft and does not take into account the possibility for the C-terminal tail of one monomer to bind in the ssDNA-binding cleft of another monomer. (B) Mechanism of the interaction of the C-terminal tail with the gp5/DNA polymerase. When the DNA-binding cleft is occupied (ssDNA is depicted in red), the negatively charged gp2.5 C-terminal tail (in red) interacts electrostatically with loop A (residues 275–285 of gp5, depicted in blue) and loop B (residues 299–314 of gp5, depicted in blue) of the TBD of the gp5/DNA polymerase. Loops A and B are easily accessible for protein–protein interactions as seen from the crystal structure of the complex of gp5/trx/primer template (omitted for clarity). The C-terminal phenylalanine of gp2.5, which is essential for the interaction with the polymerase, is not depicted because of lack of experimental data for its position.

In conclusion, it appears that the protection of the positively charged DNA-binding cleft by the electrostatic shield of the negatively charged C-terminal tail is an evolutionary conserved mechanism. This mechanism is used by abundant cellular proteins to prevent the random binding of charged molecules to their nucleic acid-binding pocket, as well as a tool for the coordination of protein–nucleic acids and protein–protein interactions within the macromolecular machinery that ensures the integrity of the genome and its replication.

Materials and Methods

Peptides and DNA. A peptide corresponding to the 26 C-terminal amino acids of T7 gp2.5 (ASKPRDEESWDEDDESEADEGDF) and a scrambled peptide with the same amino acid composition (RDDEDEDSDFKAPSEASEEWE), were synthesized by Tufts University Core Facility (Boston, MA). Single-stranded 18mer DNA was purchased from New England Biolabs, and ssDNA 23-mer oligonucleotide was purchased from Integrated DNA Technologies.

EDC/Sulfo-NHS Cross-Linking. EDC and sulfo-NHS were purchased from Pierce. EDC/sulfo-NHS is a zero-length cross-linking system that activates free carboxyl groups and cross-links them to free amino groups. A two-step cross-linking protocol was used as recommended by the manufacturer (34). The activation reaction contained 2 mM EDC, 5 mM sulfo-NHS, 50 mM potassium phosphate (pH 7), 150 mM NaCl, 0.1 mM EDTA, 10% glycerol, and 0.1 mM DTT and was performed at room temperature for 15 min. The excess cross-linking agents were quenched with 20 mM 3-mercapto-ethanol (final concentration), and the acti-
vated peptide was incubated with gp2.5 at room temperature for 20 min. The cross-linking reaction was terminated by the addition of SDS/PAGE-loading buffer and was immediately analyzed on 4–20% Tris-carboxymethylated gel (Bio-Rad).

sdDNA Cellulose-Binding Assay. sdDNA cellulose (Sigma–Aldrich) was equilibrated in 50 mM potassium phosphate (pH 7), 1 M NaCl, 0.1 mM EDTA, 0.1 mM DTT, and 10% glycerol according to the manufacturer’s recommendations. Then 500 μL of 50% slurry was loaded on a spin column and spun for 30 s at 1,000 x g. The column was washed three times with 750 μL of low-salt buffer [50 mM potassium phosphate (pH 7), 0.1 mM EDTA, 0.1 mM DTT, and 10% glycerol]. The cross-linking reaction was loaded on the DNA cellulose and incubated on ice for 30 min. The column was washed twice with 0.2 mL of low-salt buffer and eluted with 0.2 mL of the same buffer containing 1 M NaCl. The proteins in the washes and the eluate were precipitated with ice-cold TCA 10% (wt/vol) final concentration. The precipitates were washed with 70% ethanol, air-dried, and dissolved in SDS sample-loading buffer. The samples were analyzed on 4–20% SDS gels (Bio-Rad).

Expression of Isootope-Labeled Proteins for NMR. The bacterial cultures were grown on minimal medium as described previously (48). 15NH4Cl (Cambridge Isotope Laboratories) was used as the only nitrogen source to achieve uniform 15N-isotope labeling.