Structure of the θ Subunit of Escherichia coli DNA Polymerase III in Complex with the ε Subunit

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The catalytic core of Escherichia coli DNA polymerase III contains three tightly associated subunits, the αεθ, and ε subunits. The θ subunit is the smallest and least understood subunit. The three-dimensional structure of θ in a complex with an unlabeled N-terminal domain of the ε subunit, ε186, was determined by multidimensional nuclear magnetic resonance spectroscopy. The structure was refined using pseudocontact shifts that resulted from inserting a lanthanide ion (Dy3+, Er3+, or Ho3+) at the active site of ε186. The structure determination revealed a three-helix bundle fold that is similar to the solution structures of θ in a methanol-water buffer and of the bacteriophage P1 homolog, HOT, in aqueous buffer. Conserved nuclear Overhauser enhancement (NOE) patterns obtained for free and complexed θ show that most of the structure changes little upon complex formation. Discrepancies with respect to a previously published structure of free θ (Keniry et al., Protein Sci. 9:721–733, 2000) were attributed to errors in the latter structure. The present structure satisfies the pseudocontact shifts better than either the structure of θ in methanol-water buffer or the structure of HOT. This satisfies these shifts. The epitope of ε186 on θ was mapped by NOE difference spectroscopy and was found to involve helix 1 and the C-terminal part of helix 3. The pseudocontact shifts indicated that the helices of θ are located about 15 Å or farther from the lanthanide ion in the active site of ε186, in agreement with the extensive biochemical data for the θε system.

The holoenzyme DNA polymerase III (Pol III) is the main replicative polymerase in Escherichia coli (37). It is a remarkable multisubunit enzyme that is capable of extraordinary speed and fidelity of action. Pol III is composed of 10 different subunits, 7 of which act as accessory subunits for a catalytic core composed of 3 tightly bound subunits. The polymerase active site is in the large α subunit (130 kDa) of the core (33, 34). The 3'-5' proofreading exonuclease activity is located in the ε subunit (28 kDa) (48), specifically, in the N-terminal domain (ε186) (21 kDa), which also contains the binding site of the θ subunit (9 kDa). The θ subunit has no unambiguously designated function. The three subunits are arranged linearly such that the ε subunit occupies the central position, binding to both α and θ (51).

The possibility of understanding the detailed function of the Pol III holoenzyme has provoked interest in the structure of this enzyme since its discovery more than 25 years ago. Unfortunately, neither the complete holoenzyme complex nor the catalytic core has been crystallized yet. Instead, there has been considerable effort to solve the structures of individual subunits by both X-ray crystallographic and nuclear magnetic resonance (NMR) methods, and there has been substantial success recently for two of the three elements of the core. The structure of ε186 has been determined by X-ray crystallography (20) and has been modeled from NMR data (8). This structure led to a detailed understanding of the mechanism of the exonuclease activity (19, 20). NMR studies showed that the structure of the θ subunit in aqueous buffer contains three helices, a short stretch of β structure, and large segments of flexible polypeptide chains (26). Preliminary spectral data suggested that some of the poorly structured regions attain structure on binding to ε186. The refined structure of θ in a mixed alcohol-water buffer solution showed that this subunit is a three-helix bundle (39). The bacteriophage P1 homolog of θ, HOT, has the same three-helix bundle fold with minor structural variations (7). No atomic resolution structure for the α subunit or any related polymerase belonging to the PolC family is known.

The role of θ in Pol III function beyond stabilizing the structure of ε is unclear. θ is not required for the polymerase or exonuclease activities of the core or the holoenzyme (35, 50). It does, however, stimulate the exonuclease activity of ε by two- to fourfold (44, 51), enhances the interaction of ε with α (52), and stabilizes ε186 under thermal inactivation (19) and chemical denaturation (17) conditions. The biochemical evidence for a direct interaction of θ with the DNA substrate or template strand is ambiguous. HOT plays a similar role when it is substituted for θ, and there has been a report that it has an even greater stabilizing effect on ε (3). The structure of ε186 changes little when it binds to θ (6), whereas some of the poorly structured regions of θ become more ordered (26).

The θ-binding epitope on ε186 was defined using NMR chemical shift mapping experiments and was localized to a mainly hydrophobic surface encompassing strands β2 and β3, helices α1 and α2, and the N terminus of helix α7 (6). This places the θ-binding site on a face of ε186 far (approximately 15 to 20 Å) from the active site. Identification of the ε186-binding epitope on θ by similar experiments is more ambiguous.

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because chemical shift changes in θ occur for most of the molecule when it binds to ε186 (26).

One of the goals of structural genomics is to assign functions based on studies of the three-dimensional structures of proteins. With this in mind, and knowing that θ has a modest effect on the activity of ϵ, we have performed structural studies with the ε-ε186 complex to clarify the source of the effect on θ. This study is the latest in a series of studies directed at elucidating the interactions between θ, ϵ, and the DNA substrate. Here, we report the three-dimensional structure of θ when it is complexed with ε186; the results were refined using distance and angular restraints supplemented with restraints derived from pseudocontact shifts (PCS) that were induced by inserting a single lanthanide ion into the metal-binding site of ε186 (45). We compared the structure of θ in complex with ε186 with the structure of θ in mixed alcohol-water buffer and with the structure of HOT in aqueous buffer. Below we also report the results of nuclear Overhauser enhancement (NOE) difference spectroscopy experiments that defined the contact surface of θ on ε186 and compare the results with a model of the ε-ε complex that was assembled using only the PCS data (46).

MATERIALS AND METHODS

Proteins. Methods for overproduction and purification of θ and ε186 and for production and purification of the ε-ε186 complex have been described in detail previously (19, 20). 13C/15N-labeled θ and 15N-labeled ε were produced using minimal media supplemented with [1-13C]-glucose and 15N-HCl (Cambridge Isotope Laboratories) as described previously (26). Concentrations of NOESY experiment, each with a mixing time of 80 ms, were recorded at a 1H frequency of 600 MHz because chemical shift changes in the ε186 complex were measured and evaluated by using 13C and 15N heteronuclear single quantum correlation (HSQC) spectra. Well-resolved shifted cross-peaks in the 13C/H-15N-HSQC spectra of lanthamide-labeled ε-ε186 were initially assigned by inspection since they were displaced from their diamagnetic counterparts along approximately diagonal lines. The remaining cross-peaks were assigned using predicted PCS calculated from a preliminary model of θ that was derived from NOE and angular restraints. The assignments were verified by an iterative process that involved comparing the calculated PCS following structure refinement with the measured PCS derived from inspection of cross-peak positions in the diamagnetic and paramagnetic two-dimensional HSQC spectra. In addition, diamagnetic and paramagnetic 3D HNCO spectra were recorded for the ε-ε186-Dy3+ complex, and the PCS were observed in all three dimensions. This facilitated assignment of PCS that were larger in this complex than in the complexes with other lanthanide ions.

Spectral analysis and experimental restraints. Spectral weighting followed by multidimensional Fourier transformation employing linear prediction was carried out with Varian VNMRS version 6.1 software (Varian Associates, Palo Alto, CA) or MRRPipe (5) as described previously (26). After transformation, the processed data from all experiments were converted into the XEASY (1) and/or SPARKY (12) format. Peak picking, spectral analysis, and semi-automated resonance assignment were performed with XEASY, SPARKY, and CANDID (22). Three-dimensional NOESY cross-peaks were integrated using the program SPSSCAN v1.0.53 (R. W. Glaser and K. Wuthrich, http://www.mol.biol.ethz.ch/wuthrich/software/spscan). CALIBA (part of the CYANA suite of programs) (16) was used to convert NOESY cross-peak volumes into upper distance bounds. A total of 106 chemical shift index (CSI)-derived ϕ and ψ dihedral restraints from the program SHIFTY (42) were used. In addition to the NOE restraints, 48 hydrogen bond restraints (τDSET < 1.5 to 2.4 Å, τDSET = 2.4 to 3.4 Å) were included based on the CSI and other indicators of a helical structure and where a single hydrogen bond acceptor was identified in preliminary structure calculations. A total of 162 pseudocontact shifts of amide protons derived from separate experiments with 13C/15N-labeled ε-ε186-Dy3+ complex, and the PCS were observed in all three dimensions. This facilitated assignment of PCS that were larger in this complex than in the complexes with other lanthanide ions.

RESULTS

Resonance assignment and NMR spectroscopy. The optimum temperature range for acquisition of the spectra of the ε-ε186 complex was 25 to 30°C. At temperatures above 30°C, the sample aggregated, and at temperatures below 25°C, spec-
success of the 15N-HSQC spectra recorded with Dy³⁺ and in the absence of a paramagnetic ion, from which the PCS were estimated. The 15N-HSQC and 15N-separated NOESY spectra of the 13C/15N-labeled ε186 showed a chemical shift dispersion characteristic of a folded, mostly helical protein. Narrow linewidths observed for the amide protons of the residues following P70 indicated that there were mobile residues at the C terminus. Nearly complete assignment of the backbone resonances (N, HN, Cα, C') was achieved for the residues from D9 to K76, and partial assignments (N, HN) were obtained for backbone resonances for L2 to L8. The amide cross-peaks for residues L2 to L8 were broad, and their intensities in the 15N-HSQC and 15N-separated NOESY and TOCSY spectra were low. All the observable backbone amide cross-peaks (70 of an expected 71 cross-peaks) in the 15N-HSQC spectrum were assigned. Only the amide resonance of H47 could not be assigned. In contrast to the corresponding cross-peaks in the 15N-HSQC spectrum of uncomplexed ε186, in which the amide resonances in the N-terminal section of helix 1 (L8 to V18) were narrow (26), more uniform signal intensities were observed for these resonances in the ε186 complex, indicating that there was reduced mobility. Similarly, amide resonances from the C-terminal part of helix 1 (D19 to K28) and the C-terminal section of helix 3 (E54 to A62), which were either broad or not observed in uncomplexed ε (26), were easily detected and well resolved for the ε186 complex, indicating that complex formation with ε186 abolished a chemical exchange process on the millisecond timescale that broadened the resonances for free ε. In contrast, the resonances of S67 to L69 were broad for the ε-ε186 complex, although they were narrow for free ε, which may be explained by close proximity to ε186. Interestingly, DeRose et al. (7) reported low-intensity amide peaks for A22 to S24 for HOT, but the equivalent peaks for ε186-bound ε, A21 to A23, were intense. Backbone nonexchangeable 1H and 13C resonances were assigned by standard procedures using HNCA, HN(CO)CA, and HNCO spectra, which resulted in assignment of 73 Hα (96%), 70 Cα (92%), and 61 C' (80%) resonances. Most of the unassigned nonexchangeable 1H and 13C resonances were from residues near the N terminus. The absence of any intense exchange peaks between the water and the amide protons of these residues in the 15N-separated NOESY spectrum indicated that the broadening of the amide signals was due to conformational exchange rather than H exchange with the water.

Secondary structure. The NOE data were obtained from 80-ms-mixing-time 15N-separated NOESY and 13C-separated NOESY spectra at 600 and 800 MHz. The presence of medium to strong dNN and weak doN(i,i + 1) and doN(i,i + 3) NOEs and the consensus CSI (53, 54) identified three regions of helicity between Q10 and Y31 (helix 1), between A37 and R42 (helix 2), and between L48 and L66 (helix 3). Analysis using the programs TALOS (4) (Hα, Cα, C', and C' chemical shifts) and SHIFTY (42) (Hα, Cα, C', C', N, and HN chemical shifts) placed the helices at Q10 to Y31 (helix 1), between A37 and R42 (helix 2), and between L48 and L66 (helix 3).
helix 3 is amphipathic with mostly basic residues covering one face. The helical regions are broken by a sharp turn between N32 and P34, a short section of extended structure from V35 to A37, and a break between helices 2 and 3 that occurs near P45. The segment from P34 to I36 is characterized by weak dNN and strong dαN(i, i + 1) NOEs and an inversion of the CSI, indicating the presence of an extended conformation. Similarly, the pattern of strong dNN NOEs in the segment from E38 to Q44 is broken at P45 but resumes again at L48. The CSI again inverts at E46, but values characteristic of a helix residue at L48. Strong sequential dδN(i, i + 1) NOEs observed for all proline Hα resonances indicated that they are all in the trans conformation. Glycine is frequently a helix breaker (47), but all the markers for helical structure continue past G24 in the middle of helix 1. This may be due to a sequence of three alanine residues that precede G24; alanine strongly favors helix formation (47). The presence of dαN(i, i + 2) NOEs and the lack of dαN(i, i + 4) NOEs suggest that helix 1 terminates with a short stretch of a 3_10 helix.

Structure calculations. The input data for structure calculations included 1H NOEs from 15N- and 13C-separated NOESY spectra, angular data from chemical shifts, and PCS as described in Materials and Methods. The PCS data were particularly valuable as the number of long-range NOEs was relatively small (Table 1), in part because most long-range contacts involved poorly resolved side chain resonances and most of the resonances from the aromatic rings of F27, Y31, and F52 were broad.

Several rounds of structure calculations with CYANA (16) and CANDID (22) were used to refine the structure and assign cross-peaks. The final refinement was performed with the program XPLOR-NIH, with which the θ structure was refined against the complete set of distance, angular, and PCS (Dy^3+, Er^3+, and Ho^3+) restraints. After the final round of calculations, the 12 structures with the lowest energies were retained from the 200 structures calculated (Fig. 2A). The backbone conformation is well defined except for nine residues at the N terminus and 10 residues at the C terminus. The structural statistics and residual violations of the experimental restraints are shown in Table 1. Inclusion of the PCS restraints reduced the root mean square (RMS) deviation values and slightly changed the relative orientations of the helices. Refinement against the distance and angular restraints yielded backbone RMS deviations of about 1.5 Å (residues 10 to 66), whereas the RMS deviations decreased to about 1.0 Å after inclusion of the PCS restraints. No well-defined structure could be calculated for residues at the N terminus (M1 to L8) and the C terminus (P70 to K76) because of the lack of meaningful intraprotein medium- and long-range NOEs in these regions of θ. Narrow cross-peaks for residues in the region from P70 to K76 indicate that the C terminus of θ is highly mobile, whereas intermolecular NOEs between A6 of θ and residues of ε186 and conformational exchange broadening of resonances indicate that there may be some structure at the N terminus.

The quality of the structure was supported by the appearance of the Raman Chandran plot, generated by PROCHECK-NMR (28), which showed that all the residues in the generously allowed and disallowed regions are in disordered parts of the protein (mostly the C terminus), for which there were no meaningful NOE data to restrict the protein backbone to any region of conformational space.

Table 1. Structural statistics and root mean square deviations for the 12 NMR structure conformers representing the θ subunit of DNA polymerase III in complex with ε186

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
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<td>Structural restraints</td>
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<tr>
<td>Distance restraints</td>
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<td>Average restraint violations</td>
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<td>NOE violation (Å)</td>
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<tr>
<td>Largest ψ violation (°)</td>
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<tr>
<td>RMS deviations, residues 10 to 66</td>
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<tr>
<td>Backbone (Å)</td>
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<tr>
<td>Raman Chandran analysis, residues 10 to 66</td>
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<tr>
<td>Residues in the most favored regions (%)</td>
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</tr>
<tr>
<td>Residues in additionally allowed regions (%)</td>
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</tr>
<tr>
<td>Residues in generously allowed regions (%)</td>
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<tr>
<td>Residues in disallowed regions (%)</td>
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</tr>
<tr>
<td>Energetics</td>
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<tr>
<td>CHARMM energy (kcal/mol)</td>
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The overall fold of the structure (Fig. 3A) is different from the fold described previously for free θ in aqueous buffer (26) and is more similar to the fold of free θ in a mixed methanol-H2O buffer (39) and the fold of HOT in aqueous buffer (7). Although the number of long-range NOEs for θ bound to ε186 is a three-helix bundle (Fig. 2 and 3). Helix 1 is approximately antiparallel to helix 2 and parallel to helix 3, making angles of 170° and 50°, respectively. Helix 2 makes an angle of about 130° with helix 3. A hydrophobic core between the helices is populated mostly by short- and long-chain aliphatic residues. Several side chain resonances of K15 are shifted upfield, in agreement with the location of K15 between helices 1 and 3 and packed above W51. D19 and R55 are adjacent and partially protected from the solvent, and they may form a salt bridge. Several basic residues in helices 2 and 3 are oriented away from the protein core into the solution and are clustered on one side of θ. These residues are arginine residues at positions 42, 49, 53, 60, and 68.
FIG. 2. NMR-determined structures of the θ subunit in complex with ε186. (A) Stereo view of superposition of the backbone heavy atoms from the final ensemble of the 12 lowest-energy structures refined with distance, torsion angle, and pseudocontact shift restraints. Superposition was accomplished using the backbone atoms of Q10 to L66. (B and C) Calculated charge distribution mapped onto the surface of the θ subunit (residues 10 to 66). The regions with basic potential are indicated by blue, the regions with acidic potential are indicated by red, and white regions are uncharged regions of the θ surface. The mean structure is displayed, and the molecule in panel B has the same orientation as the stereo views in panels A and D, with helix 3 on the left, helix 1 in the center, and helix 2 on the right, whereas the molecule in panel C is rotated 180° about the vertical axis. (D) Stereo view of ε186-bound θ, showing some methyl-bearing side chains and the calculated position of the lanthanide ion in the active site of ε186. The side chains of methyl-bearing residues that showed substantial differential attenuation in the NOE difference experiment (>0.5) are blue. The approximate point of closest approach of the metal ion to the backbone is indicated. The coordinates are available from the PDB (PDB code 2AXD).
cause line broadening for different segments of the polypeptide chain, particularly at the interface between helix 1 and helix 3. Binding to ε186 stabilizes the θ structure rather than inducing any large changes.

HOT, which exhibits 53% sequence identity with θ (29), also folds into a three-helix bundle (7). Ribbon diagrams of ε186-bound θ, uncomplexed θ in mixed buffer (39), and uncomplexed HOT are shown in Fig. 3. Each structure has helices that are approximately the same length, and each structure has loop regions that reverse the direction of the peptide chain. The orientations of helices 1 and 3 differ by about 10° for complexed and uncomplexed θ (Fig. 3A and B) and by 20° for complexed θ and HOT (Fig. 3A and C). A DALI (23) search of the Protein Data Bank (PDB) revealed that of the proteins in the PDB, HOT is the protein most closely related to θ, with a Z score of 5.8 for the structured regions.

The structures of complexed θ and uncomplexed θ in the mixed-solvent buffer are very similar, and the greatest differences are the orientation of helix 1 with respect to the other two helices and the conformation of loop L1. These differences may be due to structural changes on complexation with ε or to the use of long-range PCS restraints. The latter better define the orientation of helices and loops, especially where there are few long-range distance restraints. Mueller et al. (39) commented on the paucity of long-range restraints in the vicinity of helix 2 and loop L2 of uncomplexed θ.

Electrostatic surface of θ. Figure 2B and C show opposing faces of the electrostatic surface of the folded core of θ (residues 10 to 66) in the complex with ε186. Figure 2C has the same orientation as the stereo views shown in Fig. 2A and D, and Fig. 2B shows the opposing face of the surface shown in Fig. 2C. The surface shown in Fig. 2B comprising mostly residues from helices 2 and 3 is highly positively charged with several arginine and lysine side chains in close proximity. The functional significance of the cluster of arginine residues oriented outward on helix 3 is unknown. There are also small pockets of hydrophobic residues at the C terminus of helix 3 and the turn structure between helices 1 and 2. Hydrophobic residues from helix 1 and helix 3 dominate the opposing face, which can be segregated into distinct regions (Fig. 2C). The surface of helix 1 is hydrophobic, resulting from the predominance of hydrophobic residues in the central part of the helix (V16 to F27). This hydrophobic surface extends to the base of the turn structure encompassing N32, M33, and P34. On either side of the central hydrophobic spine are positively and negatively charged residues.

θ-ε interface. A conventional method for defining the interface between two proteins is to map the perturbation of chemical shifts and NMR signal linewidths of one protein component caused by complexation of the other. Clustering of large changes in chemical shifts in a particular region is indicative of a potential binding surface. The θ epitope on ε186 was identified in this way (6). Figure 4A shows the changes in the chemical shift of the amide proton, amide nitrogen, backbone carbonyl, and α carbons of θ when the complex is formed. Most of the residues show significant changes in chemical shifts; the exception is the region between M33 and W51. These results suggest that the N-terminal segment of helix 1 and the C terminus of helix 3 are involved in contacts with ε186. The previously identified (26) sections of uncomplexed θ that are affected by conformational or chemical exchange phenomena include the C-terminal sections of helices 1 and 3 (D19 to Y31 and R53 to S67). Thus, it is difficult to distinguish the effects of the direct contact with ε186 from indirect effects due to a more stable secondary structure.

NOE difference spectroscopy (10) is an alternative method for mapping the binding epitope on a protein when the target protein is uniformly 13C/15N labeled and the other protein component is not labeled. We used this method to identify the side chains of methyl-bearing residues of 13C/15N-labeled θ that are at the interface with ε186. Figure 5 shows the Σ and Δ modes of part of the 1H-13C correlation map of the complex recorded with a mixing time of 300 ms. Substantial differential relaxation (>50%) was observed for the side chain resonances of A6, M13, V16, L20, A21, A23, A26, and M33 (not shown), which reside in or just precede helix 1, A62 and L66, which reside in helix 3 but face helix 1, and showed substantial differential relaxation. Figure 2D shows the locations of all of these residues in the 3D structure. Very little differential relaxation was observed for V35, I36, A37, A39, L57, A58, and V64, which reside either in the turn structure or in helix 2 or 3. A6 and L20 displayed several NOEs to the protons of unlabeled ε186 in half-filtered and double-half-filtered NOESY experiments (46). Furthermore, the methyl resonances of L20 were shifted upfield, indicating the presence of a nearby aromatic residue, which could only come from ε186 since there are no nearby aromatic rings in the θ structure. In the structure of HOT, L21 (the corresponding residue) forms part of the hydrophobic core (7), but in θ in the mixed-solvent buffer L20 is fully exposed to the solvent. The interaction interface of θ with ε186 clearly involves the hydrophobic surface of helix 1. The data are consistent with the chemical shift perturbation data (Fig. 4A), in that the residues with the smallest chemical shift perturbation show the smallest net differential relaxation effects.

An interface aligning helix 1 of θ with ε186 is also qualitatively consistent with the results of experiments with lanthanide ions at the ε186 active site, since residues in the N-terminal half of helix 1 had larger average PCS than residues in helices 2 and 3 (Fig. 4B). This indicates that helix 1 is closer to the active site of ε186 than are other regions of θ. The position
of the lanthanide ion with respect to the structure of ε is shown in Fig. 2D. The closest distance of approach of the lanthanide and helix 1 of ε, ~15 Å, is consistent with the presence of an interaction site near a hydrophobic patch on ε encompassing helix α1, strands β2 and β3, and the N terminus of helix α7 (6). Chemical shift mapping of the binding epitope of ε on ε is also suggested by the results shown in Fig. 5. Little or no change is observed in the region from V35 to R53, suggesting that this region is not in contact with ε. The resonance assignments in free ε were obtained from reference 9. (B) Plot of pseudocontact shifts of the backbone HN induced by Dy³⁺ (•), Er³⁺ (○), and Ho³⁺ (×) versus residue number.

**FIG. 4.** Changes in chemical shifts of ε upon binding to ε and ε. (A) Changes in chemical shifts of ¹H⁰ (○) and ¹⁵N[amide] (□) and ¹³Cε (•) and ¹³C′ (+) spins. There are significant changes in regions where contact between ε and ε is also suggested by the results shown in Fig. 5. Little or no change is observed in the region from V35 to R53, suggesting that this region is not in contact with ε. The resonance assignments in free ε were obtained from reference 9. (B) Plot of pseudocontact shifts of the backbone HN induced by Dy³⁺ (•), Er³⁺ (○), and Ho³⁺ (×) versus residue number.

**FIG. 5.** ¹H-¹³C correlation maps obtained by the technique of Eichmüller et al. (10), showing the differential decay of ¹H magnetization after preparation of the Σ and Δ spin modes of the ¹³C-HSQC spectra of ε-bound ¹³C/¹⁵N-labeled ε. The two spectra were recorded with an NOE mixing period of 300 ms. A reference pair of spectra obtained by using a 0-ms NOESY mixing period (not shown) was also recorded to account for differences in one-bond heteronuclear scalar coupling constants.
demonstrated that the interface between ε and θ is primarily hydrophobic (6), which is in good agreement with the results of the experiments described above. This places the ε186 epitope on θ primarily along helix 1, as shown by the hydrophobic surface in Fig. 2C. Preliminary chemical shift mapping experiments of the HOT-ε186 complex placed the interaction site near the N terminus of helix 1 and the C terminus of helix 3 (7), in good agreement with the results of the NOE difference experiments.

Notably, 13N-HSQC cross-peaks could be assigned for residues 4 to 9 preceding helix 1 in the samples with Er3+ and Ho3+. Some of these cross-peaks could also be observed for the ε-ε186-Dy3+ complex, but they were more difficult to assign. The fact that cross-peaks could be observed at all for these residues in the paramagnetic samples indicates that their amide protons must be located more than 14 Å from the paramagnetic ion (2). Cross-peaks of residues 2 and 3 were already weak in the diamagnetic complex and were therefore less likely to be observed in the paramagnetic complexes. The magnitude of the PCS observed for these residues thus indicates that they are closer to the active site of ε186 than any other residue of θ is but may not reach all the way to this site.

DISCUSSION

The small θ subunit of DNA polymerase III forms an iso-


lable complex with the catalytic N-terminal domain (ε186) of

the ε proofreading 3'-5' exonuclease subunit (19, 44). The high-resolution crystal structure of ε186 (20) has provided impetus to modeling the structure of the θ-ε186 complex, both to explain its stability and to obtain insight into the roles of θ in DNA replication. Although θ is not necessary for the exonuclease activity of ε, it has been shown to protect ε186 against thermal inactivation, as measured by the activity remaining after treatment for 10 min at an elevated temperature; the temperature at which 50% of the activity remained was about 14°C higher for ε-θ than for free ε186 (19). Free ε186 was also observed to precipitate irreversibly in a few hours under conditions used here for extensive NMR studies of the complex (at 25 or 30°C); NMR spectra of free ε186 could be obtained only at 10°C (18). Furthermore, as judged from the stability of the intact Pol III core (α-ε-θ complex) during 28,000-fold purification from plasmid-free E. coli (38), the θ-ε complex is probably extremely stable to dissociation. Direct measurement of the dissociation constant of the complex has not been reported, but an upper limit for the dissociation constant of 20 nM has been obtained by electrospray ionization-mass spectrometry (17).

Our determination of the three-dimensional structure of θ complexed with ε186 is the next step for understanding the basis for this intimate and stable protein-protein interaction, and our findings elucidated the binding interface. Remarkably, the NMR spectra obtained for θ in the 30-kDa complex with ε186 were easier to interpret than the NMR data obtained for free θ, yet the number of long-range NOEs that could be observed between the helices in θ was still small for the complex. In this situation the global PCS restraints made a significant contribution to the quality of the structure.

Overall, the structure obtained for ε186-bound θ is similar to the structure of free θ in mixed alcohol-water solvent (39) and to the structure of the close homolog HOT in aqueous buffer (7). The differences among these structures are either due to structural changes in the free and complexed proteins or due to limitations of the structure determination protocols. Since PCS restraints were used in the structure calculation, the present NMR structure of θ satisfies the PCS restraints better than the structure of either free θ (39) or HOT (7). We noted that the side chain of L21 is barely solvent exposed in the structure of HOT (7), so it would not be expected to engage in intermolecular NOEs with ε186. In contrast, the corresponding residue in θ, L20, is much more solvent exposed in the complexed and free forms of θ (39). Since the methyl groups of this leucine have intermolecular NOEs with ε186 and experience strong ring currents in the complex with ε186 but not in free θ, the two structures of θ provide plausible models of the structure in the complex. The small chemical shift changes observed for helix 2 and most of helix 3 in θ suggest that any changes in the global fold of θ upon complexation with ε186 are small.

The intermolecular NOEs observed for L20 and the NOE difference experiments (10) resulted in identification of a hydrophobic surface along helix 1, as shown in Fig. 2C, as an important part of the interaction surface with ε. The result is in agreement with the chemical shift mapping of the binding epitope of θ on ε186 that indicated that the interface between ε and θ is primarily hydrophobic (6). Electrospray ionization-mass spectrometry studies of the dissociation of ε-ε186 also suggested that the α-ε interface is hydrophobic (17). Furthermore, preliminary chemical shift mapping experiments with the HOT-ε186 complex placed the interaction site near the N terminus of helix 1 and the C terminus of helix 3 (7), in agreement with our results.

The intermolecular NOEs and reduced mobility observed for A6 show that the N-terminal segment of θ is involved in specific contacts with ε186, whereas it is highly mobile in the uncomplexed θ in aqueous (26) and mixed-solvent buffers (39). Thus, it is clear that the N-terminal segment of θ folds upon complex formation with ε186, although most interactions of this segment may be intermolecular rather than intramolecular. Although the absence of intraprotein long-range NOEs precluded structure determination for the first nine residues of θ-bound ε, PCS values and paramagnetic relaxation enhancements indicated that the N-terminal residues of θ are closer to the metal ion at the active site of ε186 than are any of the other residues of θ.

We used a novel NMR strategy (46) based on the PCS induced by inserting lanthanides at the ε186 metal-binding site to dock the NMR-derived structure of θ described above with the X-ray-derived structure of ε186 (20). The PCS-based routine for rigid docking of θ and ε186 is described in detail elsewhere (46). Briefly, the PCS detected for the amide protons of [15N]Leu- and [13C]Phe-labeled ε186 (with unlabeled θ) were used to derive the Δχ tensor of Dy3+ with respect to the crystal structure of ε186. The PCS of the amide protons of 15N-labeled θ (with unlabeled ε186) were subsequently used to position and orient the Δχ tensor with respect to the solution structure of θ described here. The common coordinate system defined in this way by the principal axes of the Δχ tensor of the metal ion positioned and oriented the structures of ε186 and θ with respect to each other. Er3+ data recorded in the same manner defined a second coordinate system that was equally
centered about the metal ion but oriented somewhat differently. The combination of Dy\(^{3+}\) and Er\(^{3+}\) data provided a unique solution for the ε186-θ complex (46). A stereo representation of the model is shown in Fig. 6. The overall structure agrees with the results of NOE difference spectroscopy, intermolecular NOEs, and chemical shift changes, placing the hydrophobic surface at the C terminus of helix 1 (Fig. 6) and part of the C terminus of helix 3 in contact with the short helix α2 in ε186. The N terminus of helix 1 of θ is approximately parallel to one of the strands of the β-sheet in ε186. The closest approach of the backbone of θ is ~15 Å from the active site of ε186, and thus the backbone of the helices of θ is too far away to have a direct role in the proofreading process. Although there is no direct interaction with the catalytically important helix 7 of ε186, θ may have a role in stabilizing the position of this helix by restraining the adjacent β strand. This stabilizing role is consistent with biological evidence showing that θ is not required for the proofreading function of ε but rather has a modest effect on the activity of ε and the fidelity of replication (52).

Exonuclease I, a structural relative of ε, has an additional C-terminal helical domain that we speculated previously may be an analog of θ (20). However, when the location of this domain in exonuclease I was compared with the location of θ on ε186, there was little overlap; the C-terminal domain of exonuclease I does not overlay the β-sheet and only partially overlays helix α2. In addition, there is very little structural homology between θ and the C-terminal domain of exonuclease I, and its function is probably unrelated (46).

The interface between θ and ε186 is large and relatively flat and is populated mostly by long-chain and aromatic side chains. The minimum contact surface between θ and ε186 as shown in Fig. 6 was calculated to be 1,400 Å\(^2\) per monomer by the program DSSP (24). The actual contact surface is almost certainly larger because the structure determination of θ did not take into account the interaction between ε186 and the N terminus of θ shown by the intermolecular NOEs between A6 of θ and ε186. This interface area corresponds to the standard-size protein interface as defined by Lo Conte et al. (30) and is typical of protein-protein complexes with nanomolar affinity (30), particularly those with hydrophobic interfaces.

Although there is no evidence that θ interacts with either single- or double-stranded DNA, it is intriguing that it has two motifs that figure prominently in DNA-binding proteins. The helix-loop-helix motif encompasses helix 1, loop 1, and helix 2 of θ, and the helix-hairpin-helix motif encompasses helix 2, loop 2, and helix 3 of θ. The helix-hairpin-helix motif has been proposed to be a nonspecific, single-stranded DNA-binding motif (9, 32), and the helix-loop-helix motif is also common in proteins that bind to DNA (21, 31). The solvent-facing surface of helix 3 of θ has several closely spaced arginine residues (Fig. 2B) that could also function as a DNA-binding region.

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