A zinc-binding site in the largest subunit of DNA-dependent RNA polymerase is involved in enzyme assembly

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All multisubunit DNA-dependent RNA polymerases (RNAP) are zinc metalloenzymes, and at least two zinc atoms are present per enzyme molecule. RNAP residues involved in zinc binding and the functional role of zinc ions in the transcription mechanism or RNAP structure are unknown. Here, we locate four cysteine residues in the *Escherichia coli* RNAP largest subunit, β' , that coordinate one of the two zinc ions tightly associated with the enzyme. In the absence of zinc, or when zinc binding is prevented by mutation, the in vitro-assembled RNAP retains the proper subunit stoichiometry but is not functional. We demonstrate that zinc acts as a molecular chaperone, converting denatured β' into a compact conformation that productively associates with other RNAP subunits. The β' residues coordinating zinc are conserved throughout eubacteria and chloroplasts, but are absent from homologs from eukaryotes and archaea. Thus, the involvement of zinc in the RNAP assembly may be a unique feature of eubacterial-type enzymes.

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DNA-dependent RNA polymerase (RNAP), alone or in complex with regulatory factors, is central to all steps of the transcription cycle, including promoter binding and melting, initiation of RNA synthesis, processive elongation, and release of RNA at terminators. Cellular RNAPs are large, multisubunit enzymes. A typical prokaryotic RNAP core contains 4–6 polypeptides with a total molecular mass of ~400 kD. Core RNAP from eukaryotes and archaea contain 7–14 subunits with a total molecular mass in excess of 500 kD. Sequence alignments of RNAP subunits reveal extensive similarities (Allison et al. 1985; Sweetser et al. 1987). Each of the two largest RNAP subunits, which are the most evolutionarily conserved, contains eight to nine colinear segments with many invariant amino acids. Low-resolution (16–35 Å) three-dimensional models of *Escherichia coli* RNAP, as well as RNAPI and RNAPII from yeast, obtained by electron crystallography, also reveal significant similarities (Darst et al. 1989, 1991; Schultz et al. 1993; Polyakov et al. 1995).

All multisubunit RNAPs are zinc metalloenzymes, and at least two zinc atoms are present per enzyme molecule (Vallee and Auld 1990). It is currently believed that each of the largest subunits of eukaryotic enzymes coordinates one zinc ion. Sequence analysis supports this view: One putative zinc-binding motif was found close

to the carboxy-terminal end of the second largest subunit from eukaryotes and archaea; another is close to the amino-terminal end of the largest subunit. Both motifs were shown to interact with radioactive zinc in blotting experiments involving denatured subunit fragments (Treich et al. 1991).

RNAP from *E. coli* is the best-studied enzyme of its class. It consists of a catalytic core of two α , one β' , and one β subunit (329, 1407, and 1342 amino acids, respectively). Binding of one of several specificity σ factors converts the core into the holoenzyme, capable of specific initiation at promoters. *E. coli* RNAP contains two zinc ions per molecule of holoenzyme (Scrutton et al. 1971). Experiments involving metal substitution and oxidative immobilization indicated that one zinc ion is associated with β' , whereas the other may be associated with either β' or β (Wu et al. 1977). However, similar experiments with the highly homologous enzyme from *Bacillus subtilis* revealed that both zinc ions are bound to β' (Halling et al. 1977).

Despite extensive research, the role of zinc atoms in RNAP structure and mechanism is not known. Addition of zinc markedly increases yields of active *E. coli* RNAP during in vitro reconstitution (Borukhov and Goldfarb 1993), and no RNAP activity can be recovered when both zinc atoms are removed (Giedroc and Coleman 1986). In contrast, removal of only one zinc ion does not prevent RNAP function (Solaiman and Wu 1985). Curiously, analysis of *E. coli* RNAP subunit sequences reveals only

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one recognizable zinc-binding motif in the β' conserved segment A. This motif corresponds to a truncated version of the putative zinc-binding motif found in the largest subunits of eukaryotic RNAP (Treich et al. 1991).

Our working hypothesis is that zinc has primarily a structural role and promotes RNAP assembly by helping bring the large subunits together in a defined orientation. This hypothesis is in accord with available structural data, showing that protein loops stabilized by zinc ions are involved in the formation of quaternary structure in other multisubunit proteins (Vallee and Auld 1990). Here, we used in vivo metal substitution, localized hydroxyl radical cleavage, and site-directed mutagenesis to locate one of the two *E. coli* RNAP zinc ions in the largest subunit, β' . We also used in vitro RNAP reconstitution to demonstrate that β' interacts with zinc in the absence of other RNAP subunits, and that this interaction is necessary for active RNAP assembly in vitro.

Results

Zinc suppresses the temperature-sensitive phenotype of *rpoC* assembly mutations

Recently, we mapped several *rpoC*(β') mutations that affect RNAP assembly or result in temperature-sensitive RNAP activity (Nedea et al. 1999). Two mutations, *R120* and *XH56*, occurred close to the evolutionarily conserved segment G of β' . Both *R120* and *XH56* changed amino acids that are strictly conserved within eubacterial RNAPs, but are absent from the β' homologs from archaea and eukaryotes (Fig. 1A). Three additional amino acids in this region, Cys-888, Cys-895, and Cys-898, are also conserved in eubacteria (Fig. 1A). Groups of three or more cysteines can coordinate structural or catalytic

zinc ions and ion clusters in proteins, and pairs of cysteines separated by two amino acids often serve as zinc-binding nuclei (Vallee and Auld 1990). We hypothesized that Cys-888, Cys-895, and Cys-898 may coordinate zinc, and that the *R120* and *XH56* mutations alter the zinc-binding site and thus interfere with RNAP assembly at the restrictive temperature (Nedea et al. 1999).

Studies of the yeast GAL4 protein showed that mutations in the zinc-binding motif could be rescued by simple addition of excess zinc in the growth medium (Johnston 1987). Accordingly, we tested if the temperature-sensitive phenotype of the *R120* and *XH56* strains could be corrected by the addition of zinc to the medium. As controls, we used *E. coli* XL1-blue (*rpoC*⁺) as well as cells carrying *rpoC*^{Ts4} (Nedea et al. 1999) and *rpoA112* (Igarashi et al. 1990) temperature-sensitive mutations. The results are presented in Figure 1B. In the absence of zinc, only XL1-blue cells grew at the restrictive (42°C) temperature, as expected. In contrast, in the presence of 100 μ M zinc in the growth medium, the temperature-sensitive phenotype of *R120* and *XH56* strains was corrected. Control cell carrying the *Ts4* and *rpoA112* mutations did not grow at 42°C even in the presence of zinc. Thus, this in vivo result supports the idea that *R120* and *XH56* affect a zinc-binding domain of β' .

Monitoring RNAP zinc-binding sites using localized radical cleavage

The Fe²⁺ ion under aerobic conditions generates reactive hydroxyl radicals, which can cleave nearby biopolymers. This process, known as the Fenton reaction, is often used for footprinting nucleic acid-protein complexes (Dixon et al. 1991). Zn²⁺ in zinc-finger proteins can be efficiently replaced by Fe²⁺ in vivo, and the resulting 'iron fingers'

Figure 1. The *R120* and *XH56* *rpoC* assembly mutations are suppressed by zinc. (A) Genetic context of the *R120* and *XH56* mutations. The bar at the top represents the 1407-amino-acid β' subunit from *E. coli*. The lettered boxes indicate evolutionarily conserved segments. *E. coli* segment G is expanded underneath and is aligned to the corresponding segments from *M. leprae* (M.L.), *T. maritima* (T.M.), *H. pylori* (H.P.), and tobacco chloroplasts (T.), as well as *S. acidocaldarius* (S.A.), and yeast RNAP I, II, and III (Yp1, Yp2, and Yp3, respectively). The *XH56* and *R120* mutations are indicated above the *E. coli* sequence; the three cysteines conserved within the eubacterial lineage are indicated below the tobacco sequence. (B) The overnight growth of the indicated *E. coli* strains on LB plates with or without 100 μ M ZnCl₂ at 30°C and 42°C.

generate free radicals that cleave both nucleic acids and polypeptide chains close to the metal-binding site (Conte et al. 1996). Earlier studies demonstrated that both *E. coli* RNAP Zn²⁺ ions could be substituted for other metal ions without affecting RNAP structure and function. RNAP purified from cells grown on minimal medium supplemented with cobalt instead of zinc contained two molar equivalents of cobalt, and no zinc and was fully functional in vitro (Chatterji and Wu 1982). Similarly, in vitro reconstitution at controlled conditions allowed the preparation of functional RNAP containing Co, Cd, or Hg ions (Solaiman and Wu 1985).

We repeated the RNAP metal biosubstitution procedure of Chatterji and Wu (1982) and purified RNAP from cells grown in zinc-depleted, iron-enriched medium. The resultant RNAP was functional and absorbed light at 400–500 nm (data not shown). In contrast, the standard RNAP preparation does not absorb light in this range. We conclude that RNAP purified from zinc-depleted, iron-enriched medium contains tightly bound iron, and we refer to this preparation as RNAP^{Fe}.

Next we used RNAP^{Fe} in localized radical cleavage reaction. Open promoter complexes were formed on a DNA fragment containing the T7 A2 promoter with either RNAP^{Fe} or control RNAP^{Zn} immobilized on Ni²⁺–NTA agarose beads through His₆-tagged β'. RNAP was then affinity labeled, using chimeric rifampicin–GTP compound and derivatized thymidine triphosphate, as described in Materials and Methods. Affinity labeling re-

sults in covalent attachment of Rif–GpCpT* to the conserved segment G of β' (Severinov et al. 1996; the bold-face type indicates radioactive phosphate; the asterisk indicates cross-linked nucleotide). The immobilized affinity-labeled complexes were washed from unincorporated reaction substrates and incubated with H₂O₂ and ascorbate, and the products were separated by SDS-PAGE and visualized by autoradiography (Fig. 2A).

Two cleavages were specific for RNAP^{Fe} (Fig. 2, lanes 7,8, sites I and II) and were thus due to the presence of tightly bound Fe²⁺. To map the iron-cleavage sites, the radio-tagged β or β' subunits were cleaved at Met residues under the 'single-hit' conditions to yield families of nested, easily identifiable fragments (Fig. 2, lanes 1,2). From the relative migration of products, the major cleavage site (site I) was mapped at the amino-terminal side of β' Met-932, which creates the smallest radioactive fragment produced under single-hit CNBr cleavage conditions. The second cleavage site (site II) was mapped close to Met-821–Met-822. We used the recombinant fragment of β', β'_{821–1407} (Severinov et al. 1996) as a marker to ensure that the assignment of single-hit bands is correct (as can be seen in Fig. 2B, the band corresponding to site II cleavage migrates just above β'_{821–1407}, which is indicated by an arrow). Thus, the site II cleavage occurred carboxy terminal to the β' conserved segment F (Fig. 2B) involved in transcription elongation and termination (Waelbaeher et al. 1994), the binding of streptolydigin in bacterial RNAP (Severinov et al. 1995; Yang and Price

Figure 2. Localized radical cleavage of wild-type RNAP. (A) RNAPs containing zinc or iron were affinity labeled in β' segment G using the T7 A2 promoter-containing DNA fragment as a template and treated with 0.03% H₂O₂ and 1 mM sodium ascorbate for times indicated. The reaction products were resolved on a 10% SDS-polyacrylamide gel and visualized by autoradiography. (Lanes 1,2) Marker lanes generated by treating the RNAP affinity labeled at β Lys-1065 (lane 1) or β' conserved segment G (lane 2) with CNBr under single-hit conditions. The mobility of the recombinant β'_{821–1407} fragment is indicated by an arrow at the right of the gel. The bar at right represents *E. coli* β'. The affinity-labeling site in segment G is shown. The three catalytic Asp residues in segment D, the four evolutionarily conserved Cys residues in segment A, as well as the cysteines close to cleavage sites I and II are shown as shaded lines. (B) Genetic context of the β' iron cleavage site II (see Fig. 1A legend for details). Cys-814 is highlighted by larger font size; an *E. coli* mutation S793F that results in streptolydigin-resistant RNAP (Severinov et al. 1995) is shown above the *E. coli* sequence. (↓) The position of natural split site in archaea.

1995), and the binding of α -amanitin in eukaryotic RNAP (Bartolomei and Corden 1987). Similar to site I, there is a cysteine residue at or very close to cleavage site II, Cys-814, that is strictly conserved within the eubacterial lineage but is absent from archeal and eukaryotic β' homologs.

Both RNAP^{Zn} and RNAP^{Fe} were cleaved with the same low efficiency near the β' conserved segment D (Fig. 2A, lanes 4,7). It has been shown that three Asp residues in segment D that normally coordinate a loosely bound Mg²⁺, which is important for RNAP catalytic activity, can also coordinate a ferrous ion (Zaychikov et al. 1996). It should be noted that the experiment of Figure 2A was performed in the absence of Mg²⁺ ions. When the cleavage reaction was repeated in the presence of 10 mM Mg²⁺, the segment D cleavage was suppressed completely (data not shown). In contrast, cleavage specific for RNAP^{Fe} was not affected by 10 mM Mg²⁺ in the reaction (not shown). Thus, the observed segment D cleavage is likely explained by the presence of residual iron in the buffer.

Localized radical cleavage experiments were also performed with RNAP^{Fe} affinity labeled at the β subunit Lys-1065 (Grachev et al. 1987). No difference in radio-tagged β cleavage between RNAP^{Fe} and RNAP^{Zn} was observed (data not shown), suggesting that the second largest subunit does not participate in the binding of Zn²⁺.

Mutational analysis of the β' zinc-binding site

The results of localized radical cleavage further support the idea that Cys-888, Cys-895, and Cys-898 are part of a zinc-binding site. Based on the identical kinetics of accumulation of the site I and site II cleavage products (data not shown), cleavage at both sites originates from the same ferrous ion, indicating that β' residues Cys-888, Cys-895, Cys-898, and Cys-814 jointly form one of RNAP zinc-binding sites. We performed site-directed mutagenesis of site I and site II cysteines of the *rpoC* gene cloned on the expression plasmid pCYB2 β' (Nedea et al. 1999). Our goal was to use mutant enzymes (1) to confirm the role of sites I and II cysteines in the coordination of zinc ion, and (2) to study the structural and functional consequences of zinc ion removal. The first mutation, 3C3A, changed Cys-888, Cys-895, and Cys-898 of site I to Ala. The second mutation, C814A, changed the only site II cysteine to alanine. Plasmids pCYB β' C814A and pCYB β' 3C3A expressing the mutant β' subunits did not complement the chromosomal *rpoC* 39°C temperature-sensitive mutation at 42°C (Nedea et al. 1999). At 30°C these plasmids had no effect on cell growth. Thus, the mutant *rpoC* genes were either temperature-sensitive or recessive lethals.

The β' subunit expressed from pCYB β' _{His} and its derivatives is His₆ tagged at its carboxyl terminus, which enables the separation of RNAP containing plasmid-borne β' from RNAP containing chromosome-encoded β' by affinity chromatography on Ni²⁺ sorbent. Our repeated attempts to purify RNAPs harboring the C814A

mutation from cells grown at 30°C using affinity chromatography failed (data not shown). We conclude that C814A completely abolishes the ability of β' to enter *E. coli* RNAP core in vivo. In contrast, affinity-purified RNAP carrying the 3C3A mutation was active in vitro (Fig. 3). Because 3C3A was engineered near the XH56 and R120 mutations that make RNAP activity temperature sensitive (XH56) or interfere with RNAP assembly in vitro (R120) (Gross et al. 1977), we asked whether RNAP^{3C3A} is active at high temperature of 42°C and whether active RNAP^{3C3A} can be obtained after in vitro denaturation/renaturation. As can be seen, the 3C3A enzyme was functional at high temperature (Fig. 3A), but it did not assemble after denaturation in vitro (Fig. 3B) and thus resembled RNAP^{R120}.

To rigorously test our interpretation that the 3C3A triple mutation disrupts an RNAP zinc-binding site we prepared iron-substituted 3C3A enzyme and performed localized radical cleavage (Fig. 3B). As expected, no cleavage at site I and site II was observed in 3C3A RNAP^{Fe}, further supporting the notion that both cleavages originate from the same iron atom. Interestingly, a major cleavage site close to one of the termini of β' was present in the 3C3A enzyme (lane 3). A well-defined putative zinc-binding motif is present in the amino-terminal segment A (amino acids 85–117) of β' (Borukhov et al. 1991). No such motif is present close to β' carboxyl terminus. Therefore, we tentatively assigned this new cleavage to the amino terminus of β' , and we mapped it close to amino acid position 115 using recombinant amino-terminally truncated β' fragments $\beta'_{88-1407}$ and $\beta'_{175-1407}$ as markers.

In vitro reconstitution of mutant enzymes in the presence of Zn²⁺ and RNAP^{WT} reconstitution in the absence of Zn²⁺ leads to inactive RNAP complex

We investigated the assembly defect caused by the C814A and 3C3A mutations using in vitro RNAP reconstitution from recombinant subunits and a Ni²⁺-coimmobilization assay to test the ability of His₆-tagged mutant β' to interact with RNAP α , β , and σ^{70} subunits. In parallel, we studied RNAP^{WT} assembly in the absence of zinc. Reconstitution mixtures were incubated with Ni²⁺-NTA agarose beads, the beads were washed and the bound material was eluted with 100 mM imidazole and analyzed by SDS-PAGE (Fig. 4). As can be seen, α , β , and σ^{70} were specifically coimmobilized with C814A and 3C3A β' subunits in the expected stoichiometry. Similarly, wild-type β' assembled into a holoenzyme-like complex both in the presence and in the absence of zinc. We conclude that C814A and 3C3A do not prevent the β' interaction with other RNAP subunits and that this interaction can occur in the absence of added zinc or when zinc binding is prevented by mutation.

The C814A, 3C3A, and the wild-type RNAP complexes assembled in the absence of zinc were transcriptionally inactive and did not interact with promoter DNA (data not shown). All three complexes exhibited

Figure 3. In vitro transcription by 3C3A RNAP. (A) RNAP^{3C3A} is temperature resistant in vitro. RNAP^{3C3A} was affinity purified from 397C cells harboring the pCYB2β^{3C3A} plasmid and used to transcribe the T7 A1 promoter-containing DNA fragment at the indicated temperatures. Reaction products were separated on a denaturing 20% polyacrylamide gel and revealed by autoradiography. (B) RNAP^{3C3A} is assembly defective in vitro. RNAP^{3C3A} and RNAP^{WT} were subjected to a denaturation/renaturation cycle by the addition of 6 M guanidine-HCl, followed by dialysis into transcription buffer. The samples were then assayed in a steady-state transcription assay, using the T7 A1 promoter as a template. Reaction products were analyzed as in A. (C) Localized radical cleavage of iron-containing RNAP^{3C3A}. RNAPs containing plasmid-borne, His₆-tagged wild-type or 3C3A β' were affinity purified from 397C *E. coli* cells grown in the presence of iron and in the absence of zinc. Affinity labeling, hydroxy-radical cleavage, and reaction-product analyses were performed as described in the legend to Fig. 2.

abnormal and similar chromatographic behavior on SEC-400 size-exclusion HPLC column: They eluted early as broad peaks. In contrast, wild-type RNAP assembled in the presence of zinc eluted later as a sharp peak (Fig. 4). Limited proteolysis with trypsin also confirmed that structural organization of reconstituted C814A and 3C3A enzyme is different than the wild-type RNAP reconstituted in the presence of zinc and is similar (or identical) to that of RNAP^{WT} reconstituted without zinc (data not shown). We conclude that in the absence of zinc, or when zinc binding is prevented by the C814A or 3C3A mutations, the specific interaction of β' with α₂β still occurs but leads to inactive complex, probably resembling the 'immature' state detected in the earlier studies (Zillig et al. 1977).

β' becomes assembly competent in the presence of zinc

The results from the previous section suggest that (1) zinc is required for proper folding of β', or (2) zinc acts at stages after β' association with α₂β to properly fold RNAP complex. To distinguish between these possibilities, we performed RNAP reconstitution from β' or α₂β renatured separately either in the presence or in the absence of zinc. The results are presented in Figure 5A. When β' and α₂β were renatured separately in the presence of zinc and combined together, active RNAP was assembled, as expected (lane 3). The yield of the assembled enzyme was about five times lower than that

obtained by standard reconstitution procedure, when α, β, and β' were renatured together in the presence of zinc (lane 1). As expected, no RNAP activity was recovered when α, β, and β' were renatured in the absence of zinc, either together or separately (lane 2; data not shown). In contrast, when β' was renatured in the presence of zinc, RNAP activity was recovered even when α₂β was assembled in the absence of zinc (lane 5). Importantly, zinc had to be present during the renaturation of β' from the fully denatured state, as addition of zinc to β' renatured in its absence did not restore the activity (lanes 6,8).

When wild-type β' was renatured individually in the absence of zinc and loaded on a gel-filtration column, it eluted as a diffuse, high-molecular-weight peak (Fig. 5B). In contrast, when β' was renatured in the presence of zinc, an additional, narrow peak with higher retention time appeared. No such narrow peak was observed when the β' subunits carrying the C814A and 3C3A mutations were renatured in the presence of zinc (Fig. 5B).

Both the diffuse peak and the narrow peak contained β', as judged by SDS-PAGE analysis (Fig. 5B). Only the material from the narrow peak, however, was able to form active RNAP when combined with α₂β and σ, as the gel shown on the bottom of Figure 5B demonstrates. From these experiments we conclude that (1) zinc is able to interact with β' in the absence of other RNAP subunits; (2) zinc converts β' into an assembly-competent, compact state; and (3) the C814A and 3C3A mutations prevent the interaction of zinc with β' and thus interfere with the assembly of functional RNAP (Fig. 5B).

Figure 4. Zn^{2+} is not important for interactions of β' with other RNAP subunits. (A) Binding of $\alpha_2\beta$ to immobilized $\text{His}_6\text{-}\beta'$ studied by the Ni^{2+} -coimmobilization assay. The α subunit was renatured with β , σ , and $\text{His}_6\text{-}\beta'^{\text{WT}}$ in the presence or in the absence of $10\ \mu\text{M}$ ZnCl_2 (lanes 1–5 and 6–10, respectively), or with $\text{His}_6\text{-}\beta'^{3\text{C}3\text{A}}$ (lanes 11–15) or $\text{His}_6\text{-}\beta'^{\text{C}814\text{A}}$ (lanes 16–20) in the presence of ZnCl_2 , loaded onto Ni^{2+} -NTA-agarose beads (L), and the unbound protein (or flowthrough) removed (F). The beads were then washed with buffer containing $5\ \text{mM}$ imidazole (W1) and $25\ \text{mM}$ imidazole (W2), and the bound proteins were eluted with buffer containing $100\ \text{mM}$ imidazole (E). The protein fractions were analyzed by SDS-PAGE. Control experiments indicated that when $\text{His}_6\text{-}\beta'$ was omitted from the reconstitution mixture, no nonspecific binding of β and α to the Ni^{2+} -NTA-agarose beads occurred (data not shown; Wang et al. 1997). (B) Abnormal chromatographic behavior of RNAP assembled in the absence of zinc. Proteins eluted from Ni^{2+} -NTA beads (lanes 5, 10, 15, 20 in A) were loaded on a SEC-400 gel-filtration column.

Discussion

The results of our work directly implicate four cysteine residues at positions 814, 888, 895, and 898 of the largest *E. coli* RNAP subunit, β' , in zinc binding and RNAP assembly. First, the temperature-sensitive phenotype of the *R120* and *XH56 rpoC* assembly mutations that occurred very close to Cys-888, Cys-895, and Cys-898 can be suppressed by the simple addition of extra zinc in the medium. Second, using RNAP containing ferrous ions instead of zinc, we can demonstrate efficient cleavage close to Cys-888, Cys-895, and Cys-898 (site I) and Cys-814 (site II). Based on the identical kinetics of accumulation of cleavage products, both site I and site II cleavages likely originate from a single ferrous ion. Finally, both site I and site II cleavages are abolished by the *3C3A* mutation, which is expected to destroy zinc binding.

The site I and site II cysteines are separated by >70 amino acids of β' primary sequence and are located in different structural modules of the β' subunit. The β' modules can be physically separated from each other without preventing RNAP assembly in vitro (Severinov et al. 1996). The binding of zinc may bring the two modules of β' together in correct orientation. Consistent with this interpretation, we were able to show that zinc converts β' into an assembly-competent and compact conformation. In vitro, β' interacts with other RNAP

subunits even in the absence of zinc or when zinc binding is prevented by mutation. Hence, the primary determinants of β' interaction with the $\alpha_2\beta$ subassembly must lie outside of the zinc-binding domain defined in this work. Zinc had no effect on $\alpha_2\beta$ formation, and no cleavage of the β subunit was detected. These results suggest that β is not involved in zinc binding, in agreement with earlier data obtained with RNAP from *B. subtilis* (Halling et al. 1977).

The *3C3A* mutation behaves as either a recessive lethal or temperature sensitive in vivo and abolishes active RNAP assembly in vitro. RNAP^{3C3A} purified from cells, however, is active, suggesting that binding of zinc in this site is not absolutely necessary for RNAP activity and assembly in vivo, at least at low temperature. Alternatively, it is also possible that other factors (e.g., chaperones) or mechanisms (e.g., cotranslational folding) may contribute to efficient RNAP assembly, even when zinc binding is prevented by mutation. The site I and site II cysteines are absent from the largest, β' -like subunits of eukaryotic and archeal RNAPs. Thus, RNAP assembly mechanisms in these organisms may differ from those used by prokaryotes. For example, small subunits of eukaryotic RNAPs interacting with the largest subunit can promote eukaryotic RNAP assembly in vivo (Gadal et al. 1999; Voutsina et al. 1999). Regardless of that fact, the β' residues responsible for zinc coordination are strictly

Figure 5. The β' subunit becomes assembly competent in the presence of zinc ions. (A) The β' subunit or a mixture of the α and β subunits (labeled $\alpha_2\beta$) were renatured separately either in the presence or in the absence of $10 \mu\text{M}$ ZnCl_2 . Proteins were then dialyzed in a buffer without zinc, combined as indicated, and incubated for 15 min at 30°C with the recombinant σ^{70} subunit to reconstitute RNAP either in the presence (lanes 6–8) or in the absence (lanes 1–5) of $10 \mu\text{M}$ ZnCl_2 . RNAP activity was then assayed using DNA fragment containing the T7 A1 promoter (see Fig. 3A legend). (Lanes 1,2) Controls, where stan-

dard RNAP reconstitution reactions containing α , β , and β' were carried out either in the presence (lane 1) or in the absence (lane 2) of zinc. (B) Chromatographic analysis of the β' subunits individually renatured in the presence or in the absence of zinc. The indicated β' subunits were individually renatured and analyzed on an SEC-400 column attached to HPLC. Fractions were collected and analyzed by 8% SDS-PAGE (top). The gel at the bottom shows the results of in vitro transcription reaction using the indicated chromatographic fractions of wild-type β' renatured in the presence of zinc. Fractions were supplemented with $\alpha_2\beta$ and σ and assayed as in A.

conserved in eubacteria and chloroplasts, suggesting that zinc plays an essential and ubiquitous role in RNAP assembly in these systems.

The *C814A* mutation appears to be more severe in vivo than *3C3A*. This result is unexpected, as both *C814A* and *3C3A* are expected to destroy the same zinc-binding site, and in vitro, recombinant β' subunits carrying either of the two mutations appear to be equally defective in RNAP assembly. Therefore, the in vivo effect of *C814A* is evidently not a simple consequence of the disruption of the zinc-binding site. We speculate that *C814A* makes RNAP proteolitically labile, thus explaining our inability to prepare active RNAP^{C814A}.

It has been shown previously that the two zinc ions bound to *E. coli* RNAP are not equal and that one, the so-called 'loosely bound' zinc ion can be removed or substituted for another metal ion under relatively mild reducing conditions without protein denaturation or loss of activity (Solaiman and Wu 1984). The *3C3A* enzyme is functional, suggesting that the zinc-binding site defined in this work is 'loose.' We attempted to prove this conjecture experimentally by performing in vitro metal (zinc for iron) substitution experiments. Unfortunately, the conditions of in vitro metal exchange also promote Fenton cleavage, and no functional RNAP could be recovered (data not shown).

Localized radical cleavage results obtained with iron-substituted *3C3A* enzyme are consistent with the idea that the universally conserved segment A also coordinates zinc. This zinc-binding site, however, appears to be inaccessible to Fenton oxidation in the wild-type enzyme. Disruption of the carboxy-terminal β' zinc-binding site by the *3C3A* mutation makes the segment A site accessible for cleavage, supporting the notion that segment A may indeed coordinate zinc. Thus, the two RNAP zinc-binding sites may interact with each other. Interestingly, genetic analysis of yeast RNAP I also suggested that segment A of the largest, β' -like subunit may interact with the putative zinc-binding motif located in the carboxyl terminus of the second largest, β -like subunit (McCusker et al. 1991). Our ongoing mutational analysis of β' segment A should clarify its role in zinc binding, RNAP structure, and mechanism.

Materials and methods

Mutagenesis of the cloned *rpoC* gene

The pCYB β ^{WT} *rpoC* expression plasmid (Nedea et al. 1999) was used to generate *rpoC* mutants. The *3C3A* mutation was generated by a single-step PCR using *3C3A*, a 71-base DNA oligonucleotide as an upstream, mutagenic primer and a downstream primer that is complementary to *rpoC* codons 1400–1407 and contains an in-frame *XhoI* site. The *3C3A* primer corresponds to *rpoC* codon positions 877–902 and introduces three Cys → Ala substitutions. In addition, it contains a *SalI* site, corresponding to the natural *rpoC* site at codon 877. The 2000-bp PCR product was cloned into pT7-blue blunt vector (Novagene), excised with *SalI* and *XhoI*, and cloned into appropriately treated pCYB β ^{WT}. After transformation in *E. coli* XL1-Blue, cells transformants were screened for the appearance of inducible β' polypeptides on

SDS–polyacrylamide gels, and the presence of mutation was confirmed by DNA sequencing.

To generate the *C814A* mutation, an 843-bp *rpoC* fragment was amplified with C814 DNA oligonucleotide as a downstream, mutagenic primer and an upstream primer corresponding to *rpoC* codons 541–548. The *C814A* oligonucleotide is complementary to *rpoC* codon positions 815–822 and introduces Cys-814 → Ala substitution. In addition, it contains a *BspHI* site, corresponding to the natural site at *rpoC* codon 819. The PCR product was cloned in pT7-blue, the 827-bp *SphI*–*BspHI* *rpoC* fragment was excised and subcloned into appropriately treated pACYC184. The resultant plasmid was treated with *BspHI*–*HindIII* and ligated with a 2500-bp *BspHI*–*HindIII* fragment of pCYB β ^{WT} containing the end of *rpoC*. Finally, a 2393-bp *BsmI*–*XhoI* fragment containing *C814A* mutation was recombined into the appropriately treated pCYB β ^{WT}. After transformation in *E. coli* XL1-Blue cells, transformants were screened for the appearance of inducible β' polypeptides with increased mobility on SDS–polyacrylamide gels, and the presence of mutation was confirmed by DNA sequencing.

RNAP purification

The β' subunits expressed from pCYB β ^{WT} or its derivatives contain a carboxy-terminal hexahistidine tag. pCYB β ^{WT}, pCYB β ^{3C3A}, and pCYB β ^{C814A} were transformed into 397C (Nedea et al. 1999) *E. coli* cells, and transformants were grown in 4 liters of Luria broth (LB) without addition of IPTG until late log phase. The cells were collected and disrupted by passing through a French press. The pellet, which contained most of the plasmid-borne β' was used in RNAP reconstitution experiments (see below). The supernatant was used for RNAP purification by a combination of polyethyleneimine (Polymix-P) precipitation, metal ion affinity chromatography, and heparin–agarose affinity chromatography. RNAP was concentrated by filtration through a C-100 concentrator (Amicon, USA) to ~1 mg/ml and stored in 50% (vol/vol) glycerol storage buffer at –20°C.

For RNAP^{Fe} purification, 397C cells transformed with pCYB β ^{WT} or pCYB β ^{3C3A} were grown in 2 liters of M9 minimal medium containing 50 μ M FeCl₃ and induced as described above. All components of the M9 medium, excluding CaCl₂, Mg₂SO₄, and FeCl₃, were treated with a Chelex-100 (Bio-Rad) ion exchanger (2 grams/liter) for 3 hr with agitation. CaCl₂, Mg₂SO₄, and FeCl₃ solutions were sterilized separately. The glassware was washed with nitric acid to remove traces of divalent ions.

Radical cleavage, affinity labeling, and transcription reactions

The standard 10- μ l cross-linking reaction contained 0.5–1.0 μ g of RNAP holoenzyme immobilized on ~5 μ l of Ni²⁺–NTA agarose (Qiagen, USA), 50 ng of the 106-bp DNA fragment containing the T7 A2 promoter (Severinov and Darst 1997), and 10 μ M Rif₆–GTP chimeric initiator molecule (Mustaev et al. 1994) in standard transcription buffer containing 40 mM Tris–HCl at pH 7.9, 40 mM KCl, and 10 mM MgCl₂. Immobilized promoter complexes were incubated for 5 min at room temperature, and the excess of Rif-G was washed with excess of transcription buffer (Kashlev et al. 1996). Reaction was left in the final volume of 15 μ l, followed by the addition of [α -³²P]CTP (3000 Ci/mmol, DuPont–NEN) to a final concentration of 0.3 μ M and 15-min incubation at 37°C. Unincorporated [α -³²P]CTP was removed by washing the beads with a buffer containing 10 mM HEPES at pH 8.0, 50 mM NaCl, and 2 mM MnCl₂. Reaction was left in 10 μ l of the buffer, and 3'-deoxy-3'-bromacetamidothymidine-5'-tri-

phosphate (Severinov et al. 1996) was added to the final concentration of 100 μ M. Reaction proceeded for 30 min at 37°C, the beads were washed with 10 mM HEPES at pH 8.0, 50 mM NaCl, and 10 mM MgCl₂, and affinity-labeled transcription complexes were immediately used for radical cleavage.

Radical cleavage was initiated by the addition of 0.03% H₂O₂ and 1 mM sodium ascorbate to the freshly prepared affinity-labeled transcription complexes. After incubation at 37°C for various times, reactions were terminated by the addition of thiourea to the final concentration of 100 mM. An equal volume of SDS-containing Laemmli loading buffer was added, and reaction products were immediately resolved by SDS-PAGE and revealed by autoradiography.

The standard steady-state transcription reaction contained in 15 μ l of 50 ng of the T7 A1 promoter-containing DNA fragment, 500 mM CpA primer, 25 μ M ATP, CTP, and GTP, and 0.3 μ M [α -³²P]UTP (3000 Ci/mmol, DuPont-NEN). Reaction proceeded for 15 min at room temperature, and the products were resolved by denaturing PAGE and revealed by autoradiography. In experiment shown in Figure 3B, 12 μ l of transcription buffer containing 25 μ g of RNAP was diluted to 50 μ l with 8 M guanidine-HCl. The denaturing agent was then dialyzed away by an overnight microdialysis against 500 ml of reconstitution buffer (Borukhov and Goldfarb 1993). Transcription was performed directly in the reconstitution mixture by the addition of the T7 A1 template and transcription substrates as described above.

RNAP reconstitution

E. coli 397C cells (Christie et al. 1996) were used as a host for RNAP subunit expression to minimize contamination with wild-type β' . Standard RNAP reconstitution was performed essentially as described by Borukhov and Goldfarb (1993). In experiments that lacked Zn²⁺, all buffers were treated with Chelex-100 as described above, and 2 grams/liter Chelex was also present in the dialysis buffer during dialysis. Inclusion bodies containing overexpressed subunits were solubilized in denaturing buffer (Borukhov and Goldfarb 1993) and incubated overnight on ice in the presence of 10 mM EDTA prior to reconstitution. When RNAP subunits and subassemblies were renatured individually, β' , or an equimolar mixture of α and β were dissolved in the denaturing buffer to the final concentration of 0.25 mg/ml and dialyzed against two changes of 30-fold excess of the reconstitution buffer (with or without 10 μ M of ZnCl₂). Renatured β' was mixed with $\alpha_2\beta$ at molar ratio 2:1, reactions were incubated for 30 min at 30°C with 1 molar equivalent of σ , followed by a steady state transcription assay performed as described above.

For chromatographic assays, individually renatured β' was precipitated with ammonium sulfate, dissolved in 50 mM Tris-HCl at pH 7.9, 0.2 M KCl, 10% glycerol, 2 mM β -mercaptoethanol, and loaded on a Superose-6 column equilibrated in the same buffer and attached to Waters Biodiscovery 626 system. Chromatographic fractions were combined with $\alpha_2\beta$ and assayed for transcription activity as described above.

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Note added in proof

Recent crystallographic analysis of RNAP core from *Thermus aquaticus* established unequivocally that cysteine residues homologous to *E. coli*. β' Cys-814, -888, -895, and -898 coordinate zinc ion (G. Zhang, E. Campbell, L. Minakhin, C. Richter, K. Severinov, and S.A. Darst, in prep.)

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A zinc-binding site in the largest subunit of DNA-dependent RNA polymerase is involved in enzyme assembly

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