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 subunits, 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
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 rpoCTs4 [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 µM zinc at 42°C, control cells carrying the Ts4 and rpoA112 mutations did not grow 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’
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 Zn$^{2+}$ 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 RNAPFe.

Next we used RNAPFe in localized radical cleavage reaction. Open promoter complexes were formed on a DNA fragment containing the T7 A2 promoter with either RNAPFe or control RNAPZn immobilized on Ni$^{2+}$–NTA agarose beads through His$_6$-tagged $\beta^\prime$. RNAP was then affinity labeled, using chimeric rifampicin–GTP compound and derivatized thymidine triphosphate, as described in Materials and Methods. Affinity labeling results in covalent attachment of Rif–GpCpT* to the conserved segment G of $\beta^\prime$ (Severinov et al. 1996; the boldface type indicates radioactive phosphate; the asterisk indicates cross-linked nucleotide). The immobilized affinity-labeled complexes were washed from unincorporated reaction substrates and incubated with H$_2$O$_2$ and ascorbate, and the products were separated by SDS-PAGE and visualized by autoradiography (Fig. 2A).

Two cleavages were specific for RNAPFe [Fig. 2, lanes 7, 8, sites I and II] and were thus due to the presence of tightly bound Fe$^{3+}$. To map the iron-cleavage sites, the radio-tagged $\beta$ or $\beta^\prime$ 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 $\beta^\prime$ 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 $\beta^\prime$, $\beta^\prime$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 $\beta^\prime$821–1407, which is indicated by an arrow]. Thus, the site II cleavage occurred carboxy terminal to the $\beta$ conserved segment F [Fig. 2B] involved in transcription elongation and termination (Waelbaecher et al. 1994), the binding of streptolydigin in bacterial RNAP (Severinov et al. 1995; Yang and Price

![Figure 2](image-url)
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 archael and eukaryotic β’ homologs.

Both RNAPZn and RNAPFe 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 RNAPFe 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 RNAPFe affinity labeled at the β subunit Lys-1065 [Grachev et al. 1987]. No difference in radiolabeled β’ cleavage between RNAPFe and RNAPZn 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 RNAP 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 β’ 88–1407 and β’ 175–1407 as markers.

**In vitro reconstitution of mutant enzymes in the presence of Zn2+ and RNAWT 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+-coimobilization assay to test the ability of Histagged mutant β’ to interact with RNAP α, β, and σ70 subunits. In parallel, we studied RNAWT 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 σ70 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
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 RNAPWT 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 $b_8$ with $\alpha_2b'$ still occurs but leads to inactive complex, probably resembling the ‘immature’ state detected in the earlier studies (Zillig et al. 1977).

$b' \text{ becomes assembly competent in the presence of zinc}$

The results from the previous section suggest that (1) zinc is required for proper folding of $b'$, or (2) zinc acts at stages after $b'$ association with $\alpha_2b$ to properly fold RNAP complex. To distinguish between these possibilities, we performed RNAP reconstitution from $b_8$ or $\alpha_2b$ renatured separately either in the presence or in the absence of zinc. The results are presented in Figure 5A. When $b'$ and $\alpha_2b$ 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 $\alpha$, $\beta$, and $b'$ were renatured together in the presence of zinc [lane 1]. As expected, no RNAP activity was recovered when $\alpha$, $\beta$, and $b'$ were renatured in the absence of zinc, either together or separately [lane 2; data not shown]. In contrast, when $b'$ was renatured in the presence of zinc, RNAP activity was recovered even when $\alpha_2b$ was assembled in the absence of zinc [lane 5]. Importantly, zinc had to be present during the renaturation of $b'$ from the fully denatured state, as addition of zinc to $b_8$ renatured in its absence did not restore the activity [lanes 6,8].

When wild-type $b'$ 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 $b'$ was renatured in the presence of zinc, an additional, narrow peak with higher retention time appeared. No such narrow peak was observed when the $b_8$ subunits carrying the C814A and 3C3A mutations were renatured in the presence of zinc (Fig. 5B).

Zinc and RNA polymerase assembly

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 pCYB2$^{3C3A}$ 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$^{3C3A}$ 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, His$_6$-tagged wild-type or 3C3A $b'$ 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.

Figure 5B
The results of our work directly implicate four cysteine residues at positions 814, 888, 895, and 898 of the largest E. coli RNAP subunit, β8, 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 β8 primary sequence and are located in different structural modules of the β8 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 αβ subassembly must lie outside of the zinc-binding domain defined in this work. Zinc had no effect on αβ 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. RNAP3C3A 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 archael 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 α2β) were renatured separately either in the presence or in the absence of 10 µM ZnCl2. Proteins were then dialyzed in a buffer without zinc, combined as indicated, and incubated for 15 min at 30°C with the recombinant σ70 subunit to reconstitute RNAP either in the presence (lanes 6–8) or in the absence (lanes 1–5) of 10 µM ZnCl2. RNAP activity was then assayed using DNA fragment containing the T7 A1 promoter (see Fig. 3A legend). (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 α2β 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, suggesting 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<sup>WT</sup> rpoC expression plasmid (Neda 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 an downstream primer that is complementary to nucleotide as an upstream, mutagenic primer and a downstream primer corresponding to rpoC codons 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 BspH–BspHI rpoC fragment was excised and subcloned into appropriately treated pACYC184. The resultant plasmid was treated with BspH–HindIII and ligated with a 2500-bp BspH–HindIII fragment of pCYB<sup>WT</sup> containing the end of rpoC. Finally, a 2393-bp BsmI–XhoI fragment containing C814A mutation was recloned into the appropriately treated pCYB<sup>WT</sup>. 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<sup>WT</sup> or its derivatives contain a carboxy-terminal hexahistidine tag. pCYB<sup>WT</sup>, pCYB<sup>3C3A</sup>, and pCYB<sup>C814A</sup> were transformed into 397C (Neda 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 [Polymin-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<sup>Fe</sup> purification, 397C cells transformed with pCYB<sup>WT</sup> or pCYB<sup>3C3A</sup> or pCYB<sup>C814A</sup> were grown in 2 liters of M9 minimal medium containing 50 μM FeCl<sub>3</sub> and induced as described above. All components of the M9 medium, excluding CaCl<sub>2</sub>, Mg<sub>2</sub>SO<sub>4</sub>, and FeCl<sub>3</sub> solutions were sterilized separately. The supernatant was treated with a Chelex-100 (Bio-Rad) ion exchanger [2 grams/liter] for 3 hr with agitation. CaCl<sub>2</sub>, Mg<sub>2</sub>SO<sub>4</sub>, and FeCl<sub>3</sub> 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-μl cross-linking reaction contained 0.5–1.0 μg of RNAP holoenzyme immobilized on ~5 μl of Ni<sup>2+</sup>–NTA agarose (Qiagen, USA), 50 ng of the 106-bp DNA fragment containing the T7 A2 promoter [Severinov and Darst 1997], and 10 μM Rif<sup>−</sup>–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 MgCl<sub>2</sub>. Immobilized promoter complexes were incubated for 5 min at room temperature, and the excess of Rif<sup>−</sup> was washed with excess of transcription buffer [Kashlev et al. 1996]. Reaction was left in the final volume of 15 μl, followed by the addition of [α-<sup>32</sup>P]CTP (3000 Ci/mmol, DuPont–NEN) to a final concentration of 0.3 μM and 15-min incubation at 37°C. Unincorporated [α-<sup>32</sup>P]CTP was removed by washing the beads with a buffer containing 10 mM HEPES at pH 8.0, 50 mM NaCl, and 2 mM MnCl<sub>2</sub>. Reaction was left in 10 μl of the buffer, and 3′-deoxy- 3′-bromacetamidothymidine– 5′-tri-
phosphate (Severinov et al. 1996) was added to the final concentration of 100 µM. 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 µl of 50 ng of the T7 A1 promoter-containing DNA fragment, 500 mM CpA primer, 25 µM ATP, CTP, and GTP, and 0.3 µM [α-32P]UTP (3000 Ci/m mole, 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 µl of transcription buffer containing 25 µg of RNAP was diluted to 50 µl 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 µM of ZnCl₂). Renatured β’ was mixed with αβ 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 sulfoxide, 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 αβ 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 -896 coordinate zinc ion (G. Zhang, E. Campbell, L. Minakhin, C. Richter, K. Severinov, and S.A. Darst, in prep.)

**References**


**Zinc and RNA polymerase assembly**

Note added in proof


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

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