Francisella RNA polymerase contains a heterodimer of non-identical α subunits

Damir Mukhamedyarov1, Kira S Makarova2, Konstantin Severinov1,3 and Konstantin Kuznedelov1*

Abstract

Background: All sequenced genomes of representatives of the Francisella genus contain two rpoA genes, which encode non-identical RNA polymerase (RNAP) subunits, α1 and α2. In all other bacteria studied to date, a dimer of identical α subunits initiates the assembly of the catalytically proficient RNAP core (subunit composition αββ′). Based on an observation that both α1 and α2 are incorporated into Francisella RNAP, Charity et al. (2007) previously suggested that up to four different species of RNAP core enzyme might form in the same Francisella cell.

Results: By in vitro assembly from fully denatured state, we determined that both Francisella α subunits are required for efficient dimerization; no homodimer formation was detected. Bacterial two-hybrid system analysis likewise indicated strong interactions between the α1 and α2 N-terminal domains (NTDs, responsible for dimerization). NTDs of α2 did not interact detectably, while weak interaction between α1 NTDs was observed. This weak homotypic interaction may explain low-level transcription activity observed in in vitro RNAP reconstitution reactions containing Francisella large subunits (β′, β) and α1. No activity was observed with RNAP reconstitution reactions containing α2, while robust transcription activity was detected in reactions containing α1 and α2. Phylogenetic analysis based on RpoA resulted in a tree compatible with standard bacterial taxonomy with both Francisella RpoA branches positioned within γ-proteobacteria. The observed phylogeny and analysis of constrained trees are compatible with Francisella lineage-specific rpoA duplication followed by acceleration of evolutionary rate and subfunctionalization.

Conclusions: The results strongly suggest that most Francisella RNAP contains α heterodimer with a minor subfraction possibly containing α1 homodimer. Comparative sequence analysis suggests that this heterodimer is oriented, in a sense that only one monomer, α1, interacts with the β subunit during the αβ RNAP subassembly formation. Most likely the two rpoA copies in Francisella have emerged through a lineage-specific duplication followed by subfunctionalization of interacting paralogs.

Background

Bioinformatics analysis reveals that two paralogous rpoA genes, each encoding non-identical proteins homologous to bacterial RNA polymerase (RNAP) α subunits, are present in the genome of Francisella tularensis [1]. The bacterial RNAP core enzyme has subunit composition αββ′. Variations including fusion of the largest subunits, β and β′, in Helicobacter and Wolinella genera [2,3], and split the largest subunit in some cyanobacteria [4] have been reported, but overall, the subunit composition of RNAP core is conserved. The α subunit homodimer initiates bacterial RNAP assembly. The α subunit monomers dimerize through their N-terminal domain (NTD) [5,6]. The C-terminal domain (CTD) is connected to NTD through a flexible tether [7]. The αCTD is not required for assembly but is involved in transcriptional regulation [8-10]. The αNTD homodimer provides a platform for interaction with the two large RNAP subunits [11,12]. Determinants in α important for interactions with β and β′ subunits have been localized by mutagenesis and hydroxyl-radical footprinting studies [5-8,13-15]. Substitutions at positions 45 and 48 of Escherichia coli α subunit completely (R45A) or

* Correspondence: kuznedelov@waksman.rutgers.edu
1 Department of Biochemistry and Molecular Biology and Waksman Institute of Microbiology, Rutgers, the State University of New Jersey, Piscataway, NJ 08854, USA
Full list of author information is available at the end of the article

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RNAP containing oriented $\alpha_2$ heterodimers have been prepared both in vitro, by reconstitution from recombinant subunits, and in vivo, by co-expression of genes for recombinant subunits, by using one $\alpha$ subunit lacking the R45A substitution and one $\alpha$ subunit having the R45A substitution [18,19]. Functional analysis of RNAP containing oriented $\alpha$ heterodimers confirmed that asymmetrical arrangement of $\alpha$ leads to non-identical functions of each monomer in transcription regulation [18,19].

RNAP core enzymes from archaea and eukaryotes contain homologs of each of the bacterial RNAP core subunits. However, rather than having two identical $\alpha$ subunit homologs, they contain two different $\alpha$-like polypeptides (RPB3 and RPB11 in the case of eukaryotic RNAP II) that form a heterodimer, which serves as a platform for RNAP assembly [20].

The presence of two different genes ($rpoA1$ and $rpoA2$) in the genome of Francisella suggests that up to four RNAP core enzymes differing in subunit composition could be present in the cells: two enzymes containing $\alpha$ homodimers, ($\alpha_1$)$_2$ and ($\alpha_2$)$_2$, and two enzymes containing $\alpha$ heterodimers, ($\alpha_1\alpha_2$)$_2$ and ($\alpha_2\alpha_1$)$_2$ [1]. The heterodimers could differ from one another with respect to which $\alpha$ interacts with the $\beta$ subunit of RNAP and which $\alpha$ interacts with $\beta'$ [18,19]. Promoter recognition properties of RNAP holoenzymes formed from these different core enzyme molecules may differ, since CTD of $\alpha_1$ and $\alpha_2$ may be capable of different protein-DNA and protein-protein interactions during transcription initiation [18,19]. Further, if holoenzymes containing RNAP core enzymes of different composition indeed respond differently to transcription factors and elements, then F. tularensis may regulate the spectrum of expressed genes by altering the relative ratio of core enzymes with different $\alpha$ subunit composition, which would be a novel paradigm of transcription regulation in bacteria.

Evidence of that both $\alpha_1$ and $\alpha_2$ subunits are incorporated into F. tularensis RNAP has been reported earlier by Charity et al. [1]. These authors demonstrated that RNAP affinity purified from F. tularensis strain expressing the $\beta'$ subunit with fused TAP-tag contained both $\alpha_1$ and $\alpha_2$. These experiments clearly show that both rpoA genes are active and their products are components of RNAP but do not inform about the actual subunit composition of F. tularensis RNAP. Based on predicted dimerization determinants in other bacteria [21], Charity et al. hypothesized that $\alpha_1$ and $\alpha_2$ might exclusively form either homodimers or heterodimers [1]. In the present study, we describe the results of in vitro analysis of assembly of RNAP from F. tularensis subspecies novicida. Our results indicate that RNAP core containing an $\alpha$ heterodimer is the main, perhaps the only, species of RNAP in this organism. We further present results of phylogenetic analysis that provide a plausible scenario for the appearance of two paralogous rpoA genes in the Francisella lineage.

**Results**

*F. tularensis* $\alpha$ heterodimer but not homodimers efficiently assembles in vitro

To experimentally address the ability of F. tularensis RNAP $\alpha$ subunits to form homo- and heterodimers, we investigated the ability of recombinant F. tularensis $\alpha$ subunit proteins with C-terminal His$_6$-tags to pull down untagged counterparts during ion metal affinity chromatography. As shown previously, F. tularensis RNAP $\alpha$ subunits have different electrophoretic mobilities, with $\alpha_1$ migrating significantly faster than $\alpha_2$ [1]. In addition, His$_6$-tags alter electrophoretic mobility of both $\alpha_1$ and $\alpha_2$ enough to separate tagged and untagged $\alpha$ subunits of the same kind (Figure 1).

Therefore, because each of the four proteins used in the pull-down assay has a characteristic electrophoretic mobility, it is possible to detect the efficiency of both hetero- and homodimer formation. Various pairwise combinations of $\alpha$ subunits were mixed at denaturing conditions (6 M guanidinium chloride), the denaturing reagent was removed by dialysis at conditions favouring bacterial RNAP assembly from isolated subunits [22], and reconstitution reactions (labelled “L” on Figure 1) were loaded on Ni$^{2+}$-affinity columns. Flow-through (F) was collected and retained protein was eluted (E) with different concentrations of imidazole in the buffer. Aliquots of each fraction were next analyzed by SDS-PAGE. As can be seen from Figure 1 (top panel), no co-immobilization of untagged $\alpha$ subunit in reactions that contained tagged and untagged versions of subunit of the same kind was detected. In contrast, heterodimers were readily detected when either $\alpha_1$His$_6$ or $\alpha_2$His$_6$ were used as “baits” for co-immobilization of, respectively, $\alpha_2$ or $\alpha_1$ (Figure 1, bottom panel). We conclude that F. tularensis RNAP $\alpha$ subunits do not appreciably form homodimers, at least at the conditions of in vitro RNAP assembly.

$\alpha$1NTD and $\alpha$2NTD efficiently interact in bacterial 2-hybrid system

RNAP $\alpha$ subunit is a two-domain protein, with its N-terminal domain being primarily responsible for dimerization and interaction with large RNAP subunits, while the C-terminal domain, CTD, which is connected to NTD through a flexible linker, is primarily responsible for
interactions with transcription factors and DNA upstream of the -35 promoter element [23]. Weak dimerization of isolated αCTD has been reported and may be of regulatory significance [7]. To independently study dimerization of various domains of F. tularensis α subunits, we used the bacterial two-hybrid system [24]. Eight two-hybrid plasmids expressing bait and prey fusions of each α domain were constructed and 16 pairwise combinations were tested. The results are presented in Table 1. As can be seen, in agreement with in vitro co-immobilization data, strong interactions between αNTDs of different kinds were detected. αCTDs did not appreciably interact with each other or with αNTDs. The level of homotypic interaction between α1NTD was above the background, potentially indicating formation of α1NTD homodimer, while the level of α2 homodimer formation was at the background level.

### Formation of the α2β subassembly in vitro
RNAP assembly follows a conserved pathway, whereby the β subunit interacts with the α dimer, leading to the formation of α2β - a stable intermediate of RNAP assembly that can be observed both in vivo and in vitro [11,16]. We performed in vitro RNAP assembly using His6-tagged F. tularensis α subunits and untagged F. tularensis β subunit.

### Table 1

| Fraction | L | F | E | L | F | E |
|----------|---|---|---|---|---|---|
| Imidazole (mM) | 1 | 1 | 20 | 100 | 1 | 1 | 20 | 100 |

#### Figure 1

**In vitro assembly of Francisella α homo- and heterodimers.** Reactions containing indicated proteins were combined at denaturing conditions and, following dialysis into a buffer favoring RNAP assembly, were fractionated using Ni2+-affinity chromatography. Coomassie-stained SDS gels are presented. "L" - load, "F" - flow-through, "E" - elution with buffers with indicated concentration of imidazole.

Eight two-hybrid plasmids expressing bait (1st column) and prey (1st row) fusions of each α subunit domain were constructed and 16 pairwise combinations were tested in a reporter strain by measuring β-galactosidase activity (in Miller units). Each combination was tested at least three times independently. Mean and standard deviation values are presented. Three kinds of measurements were taken to determine background levels of β-galactosidase activity, which was found to be (in Miller units) 111 ± 26 in host reporter cells with no plasmids, 110 ± 9 in cells transformed with pBRαLN-α1NTD only, and 50 ± 7 in cells transformed with pAClC-α1NTD only.
recombinant *F. tularensis* β. The results indicated that β was most efficiently immobilized when both α subunits were present in the assembly reaction (Figure 2A, lane 12). Only trace amounts of β were co-immobilized in reactions containing α2 (Figure 2A, lane 8) and thus likely represented non-specific binding (note that an excess of α2 was used in this reaction). The amount of β co-immobilized in reactions containing α1 (lane 4) was higher than the background but clearly less than that observed in reactions containing both α subunits. We conclude from these experiments that β interacts most efficiently with α heterodimer. Detected interaction between β and α1 can proceed through α1 monomer or, alternatively, the β subunit may stimulate formation of the α1 homodimer.

**In vitro transcription by recombinant *F. tularensis* RNAP**

To validate data obtained using two-hybrid analysis and α dimer/αβ RNAP subassembly *in vitro* reconstitution, *in vitro* RNAP assembly and transcription experiments were performed. Three *in vitro* RNAP assembly reactions contained recombinant *F. tularensis* β and β' subunits and α1, α2, or both α1 and α2 (the ω subunit was omitted from assembly reactions as it is not essential for

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**Figure 2** *In vitro* assembly of *Francisella* αβ RNAP subassembly. A. Reactions were assembled and analyzed as described in Figure 1 legend. ‘W’ -wash with excess of loading buffer. Proteins were eluted with a buffer containing 100 mM imidazole. B. Sequence alignment of the α subunit segment involved in dimerization and interaction with the β subunit. A segment of *E. coli* α subunit (amino acids 1-59) is shown at the top (single-letter amino acid code). Corresponding sequences from *Thermus aquaticus* (Taq), *Thermus thermophilus* (Tth), and *Francisella tularensis* RpoA variants are aligned below. Dots indicate identities, hyphens - gaps. Amino acids highlighted in red form a cluster important for α homodimer formation in *E. coli*. Amino acid highlighted in blue is responsible for the interaction with β.
RNAP basic function [25,26]). Assembled RNAP reactions were passed through a gel-filtration column, fractions that eluted at retention times expected for RNAP core elution were collected and tested for transcription activity on a nucleic acid scaffold shown in Figure 3A. Nucleic acid scaffolds mimic the conformation of nucleic acids in transcription elongation complexes. RNAP complexes with nucleic acids scaffolds are catalytically active and serve as a convenient tool to study transcription elongation properties of the enzyme [27]. Reactions were combined with NTP, and elongation of radioactively labelled 8-nt RNA component of the scaffold ("RNA₈") followed. The results are presented in Figure 3B. As can be seen, most efficient elongation of the RNA primer was observed in fractions obtained from RNAP assembly reaction containing both α subunits. Fractions of RNAP assembly reaction that contained α2 only were completely inactive. Fractions of RNAP assembly reaction containing α1 only demonstrated low but detectable transcription activity. We therefore conclude that F. tularensis RNAP assembles efficiently when both kinds of α subunits are present; α2 alone is unable to promote RNAP assembly; α1 alone supports RNAP assembly, albeit with low efficiency, possibly due to low level of α1 homodimer formation.

**Evolution of RpoA in Francisella**

To gain insight into the evolution of two paralogs of RpoA in Francisella, we retrieved all RpoA sequences in all of the 1055 completely sequenced bacterial genomes available in the RefSeq database. We found that in addition to Francisella, two rpoA genes (rpoA1 and rpoA2) are present in several other genomes, namely in three Chloroflexus species (C. aggregans DSM 9485; C. aurantiacus J-10-ß; C. sp. Y-400-ß), in Streptomyces avermitilis MA-4680, in Psychromonas ingrahamii 37, and in Leptospira borgpetersenii serovar Hardjo bovis L550 (see also Additional file 1). In the latter two cases the two rpoA copies are identical and are apparently the result of very recent genome segment duplications that also include a number of other genes.

In order to address an alternative possibility, that one of the RpoA paralogs in Francisella could have been horizontally transferred from a distant bacterial (other than γ-proteobacteria to which the Francisella genus belongs) lineage instead of arising through gene duplications listed above (Figure 4A). The resulting tree is generally very well compatible with bacterial taxonomy, which is not surprising considering the fact that RNAP subunits are among the best phylogenetic markers [28-30]. The position of both RpoA branches corresponding to Francisella within γ-proteobacteria is confidently supported by bootstrap analysis (bootstrap probability of 0.93). Thus, it is unlikely that any of the Francisella rpoA genes were transferred from outside of the γ-proteobacterial lineage. Branches leading to both Francisella RpoA proteins are extremely long, which might cause an artefact of the long-branch attraction, making the Francisella RpoA positioning unreliable. To test hypotheses for an alternative position of Francisella RpoA branches, we used RAxML [31] program to reconstruct a phylogenetic tree for γ-proteobacteria with β-proteobacteria outgroup (Figure 4B), made constrained trees and compared the maximum likelihood values for the best tree (Figure 4B) and constrained trees. The first constrained tree was designed to test a hypothesis of monophyly of two RpoA paralogs of Francisella; the second tree was designed to test a hypothesis of monophyly of both Francisella RpoA and of homologs from Coxiella, Legionella, and Thiomicrospira - species that are the closest taxonomic relatives of Francisella (Additional file 2). The analysis showed that none of the hypotheses could be rejected, suggesting that the positioning of RpoA at the root of γ-proteobacteria could be explained by long-branch attraction.

![RNA₈](5'-ACATCGCTATATTAAAGCTGTGCTGG-3' T)

**Figure 3** Transcription activity of *in vitro* assembled Francisella RNAP core enzymes. A. The structure of nucleic acid scaffold used to test assembled enzymes activities is schematically shown. B. Superose 6 fractions obtained after separation of *in vitro* RNAP assembly reactions containing indicated subunits were combined with nucleic acid scaffold containing a radioactively labelled RNA primer and reactions were supplemented with NTP. Reaction products were analyzed by denaturing PAGE. An autoradiograph is presented.
Figure 4 Phylogenetic trees of the bacterial RpoA. A set of 368 RpoA sequences from 355 representative genomes was aligned using MUSCLE program. The Maximum likelihood tree with bootstrap values applied was built for 246 informative aligned positions using FastTree program. B set of 84 RpoA sequences from 76 representative genomes of γ- and β-proteobacteria was aligned using MUSCLE program. The Maximum likelihood tree with bootstrap values applied was built for 296 aligned informative positions using RAxML program. The evolutionary model for tree reconstruction (WAG with gamma-distributed evolutionary rates) has been selected using ProtTest program. Colour code: light blue - RpoA1 subunits encoded in ribosomal operons, the default location for all bacteria; light green - RpoA2 subunits encoded elsewhere in the genome. Dashed arrows point to subtrees with two other (non-Francisella) instances of RpoA duplication: Chloroflexus and Streptomyces avermitilis. Each terminal tree node is labelled with GenBank Identifier (GI) number, five-letter taxonomy code and full systematic name of an organism. The taxonomy code is the following: Gamma - γ-proteobacteria; Beta - β-proteobacteria; Alpha - α-proteobacteria; Clost - Clostridia; Dicty - Dictyoglomi; Deino - Deinococcus/Thermus group; Syner - Synergistetes; Nitro - Nitrospira; Therm - Thermomicrobia; Dehal - Dehalococcoidetes; Fibro - Fibrobacteres/Acidobacteria group; Gemma - Gemmatimonadetes; uncla - unclassified. Compressed branches (black triangles) are labelled according to taxonomy. Branches leading to γ-proteobacteria are red. The shaded clade indicates close relatives of Francisella according to taxonomy based on 16S rRNA and concatenated ribosomal proteins phylogeny.
Discussion

Representatives of the bacterial genus *Francisella* are unusual with respect to RNAP composition, in that they contain two different α subunits encoded by two paralogous genes *rpoA1* and *rpoA2*. The presence of two different α subunits in affinity-purified RNAP preparations [1] suggested that as many as four different species of RNAP core enzyme could be present in the single cell. Here, we studied *Francisella* RNAP by means of *in vitro* assembly. Our results show that *Francisella* α heterodimer (*αα2*) efficiently assembles *in vitro* from fully denatured state and homodimers are not detected. Bacterial two-hybrid analysis indicates that in addition to efficient α heterodimer assembly, some dimerization of α1 may also occur. Thus, the efficiency of α dimerization is a clear major factor that should affect subunit composition of *Francisella* RNAP. As was determined from crystal structure analysis, the main structural elements of the α dimer interface of *E. coli* are two α-helices, H1 and H3, orthogonally oriented to each other [21]. These helices from one monomer participate in a coiled-coil-like interaction with their counterparts in the other monomer. Within these helices, Kannan et al. [32] identified a cluster of amino acids stabilizing interactions at the *E. coli* α dimer interface, with residues 35F, 38T, and 39L emanating from one α monomer and residues 46I, 50S, and 227Q from another. Of particular interest are three of them, namely, 35F, 38T, and 46I, point mutations at which partially (*α-T38A*) [16] or completely (*α-F35A, α-I46S*) [32] prevented the dimer formation. As can be seen from Figure 2B, amino acids at these positions are conserved (identical) between *E. coli* and *Thermus*, while amino acids in many of the corresponding positions in both α1 and α2 subunits from *Francisella* differ. Since these positions are critical for the dimer formation, it is reasonable to assume that some amino acids at these positions of *Francisella* α subunits, for example, α1-36M, α1-39I, α2-33V, and α2-47T, may be unfavourable to the assembly of homodimers. However, in the absence of crystal structure or systematic mutagenesis data, it is currently not possible to identify structural reasons for hetero- and homodimerization of *Francisella* α subunits.

During RNAP assembly in organisms where α subunit forms a homodimer, the β subunit is free to interact with either α monomer to form the α1β subassembly. The situation must be different in the case of *F. tularensis*, where α heterodimers form preferentially. In *E. coli*, an evolutionary conserved α subunit residue Arg45 is critical for β subunit interaction with the α dimer [16,18,19]. In *F. tularensis*, the corresponding position in α1 contains an arginine (Arg46), while in α2 this position is occupied by glutamine (Gln45; Figure 2B). Thus, it appears that β subunit in *F. tularensis* will be specifically interacting with the α heterodimer through α1. That α1 contains determinants for interactions with β also follows from results of the α1β subassembly reconstitution and *in vitro* transcription data, since *Francisella* RNAP containing α1 homodimer is functional and formation of an (α1)2β intermediate can be detected *in vitro*, albeit with low efficiency. The latter result suggests that β may stimulate α1 dimerization. An alternative possibility would be β interacting with one α1 monomer, followed by association with another α1 and β’, or the α1β’ complex. Be that as it may, our data suggest that bacteria of *Francisella* genus produce a major form of RNAP containing an oriented α1α2 heterodimer, and a minor form containing α1 homodimer.

As shown earlier by hydroxyl-radical-mediated proteolysis [14], the segments of *E. coli* α most strongly protected by β correspond to amino acids 30-55 and 65-75, and the segments of α most strongly protected by β’ correspond to amino acids 175-185 and 195-210. Single alanine substitutions of *E. coli* α Lys86 and Val173 and Two-amino-acid insertions at positions 180 and 200 of *E. coli* α cause defects in β’ binding without affecting the α1β assembly formation [16,17]. To evaluate the ability of *Francisella* α1 and α2 subunits to interact with the β’ subunit, we compared sequences of *E. coli* α subunit involved in interaction with β’ [14,21] to those in *F. tularensis* α1 and α2 subunits (Additional file 1). The results reveal that a lysine at a position corresponding to *E. coli* α position 86 is present in both α polypeptides from *Francisella*, while amino acids corresponding to *E. coli* α Val173 are, respectively, a valine and a leucine in α1 and α2. Similarly, the site of one two-amino-acid insertion that destroys β’ interaction with α2β in *E. coli*, Val180, has as its counterpart a valine in α2 and an isoleucine in α1 of *Francisella*. These conservative changes are unlikely to affect the efficiency of β’ binding by the α polypeptides. Interestingly, the site of the residue at the site of the second insertion affecting β’ interaction with α2β in *E. coli*, Lys200, is conserved in *Francisella* α2 and α subunits from other bacteria (see Additional file 1), but is substituted with threonine in α1. The results thus implies that *Francisella* β’ interacts with α2. Further experiments will be needed to prove this conjecture.

Our phylogenetic analysis indicates that none of the *Francisella rpoA* genes was transferred from outside of the γ-proteobacterial lineage. In fact, both genes most
likely emerged through duplication of an ancestral single gene followed by acceleration of evolutionary rate of both paralogs. Acceleration of rpoA evolutionary rate after the duplication apparently was accompanied by subfunctionalization of Francisella α subunits, ultimately leading to accumulation of substitutions in residues responsible for homodimerization and involved in the β and β′ subunit interaction.

Similar events, albeit on much longer time intervals, must have led to formation of two very different α-like subunits in eukaryotes and archaea. The large α-like subunit (RPB3 in eukaryal RNAP II, AC40 in RNAP I and RNAP III, Rpo3 (also known as RpoD) in archaea) heterodimerizes with its much smaller counterpart (RPB11 in eukaryal RNAP II, AC19 in RNAP I and RNAP III, Rpo11 (also known as RpoL) in archaea) [33-36]. Crystallographic and functional analyses indicate that large α homolog makes interaction with the second-largest (β-like) subunit through a surface that contains residue homologous to E. coli α Arg45 [35,37], and is thus formally similar to Francisella α1. The smaller α homolog of eukaryal and archaeal RNAP thus corresponds to Francisella α2. One should not take this analogy to far though, since in eukaryotes and archaea, the α heterodimer is not sufficient for recruitment of large RNAP subunits in the complex. Eukaryal RPB10 and RPB12 and their archaeal homologs Rpo10 (also known as RpoN) and Rpo12 (also known as RpoP) form a stable complex with all four polypeptides playing an essential role in assembly and stability of the RNAP complex [36,38,39].

Evolution of the rpoA duplication presented here is one of the best demonstrations in support of the Lynch’s subfunctionalization scenario where both copies are subject to relaxed selection and acceleration of the evolutionary rates but rarely develop a new or specia-

Conclusions

The data presented here support the following conclusions: (1) only Francisella α-heterodimer (α1α2) can be efficiently assembled in vitro; (2) strong direct interactions between α1NTD and α2NTD only have been detected in the bacterial two-hybrid system; (3) β interacts more efficiently when both of α1 and α2 presented in the reconstitution mix; (4) interaction between α1 and β subunits was observed to be stronger than interaction between α2 and β; (5) based on phylogenetic analysis, two rpoA copies in Francisella most likely must have emerged through a duplication followed by acceleration of the evolutionary rates of both paralogs.

Methods

Bacterial strains

E. coli NovaBlue Singles competent cells (Novagen) were used for initial cloning and plasmid propagation. E. coli BL21 (DE3) cells were used for protein overproduction. Reporter E. coli strain FW102 F’Ori2-62 [41] was used for bacterial two-hybrid experiments.

Cloning and expression

Francisella tularensis novicida genomic DNA has been provided by Dr. Michael Ibba (Ohio State University). Primers for PCR amplification of rpo genes were designed using Francisella tularensis subsp. novicida (FTN) strain U112 genome sequence data [GenBank: NC_008601]. The primers allowed cloning of amplified FTN rpo genes in pET series E. coli expression plasmids between the Ncol and EcoRI (or Xhol to express C-terminally hexahistidine-tagged α subunits) restriction sites. The plasmids pET28-FtnA1His, pET28-FtnA2His, pET28-FtnA1, pET28-FtnA2, and pET28-FtnB, pET30-FtnC overexpressing, respectively, C-terminally hexahistidine-tagged α1Hisα, α2Hisα, and untagged α1, α2, and β, and N-terminally hexahistidine-tagged β′ subunits, were constructed using routine cloning methods and verified by sequencing of entire rpo portions for each plasmid. BL21 (DE3) cells harbouring the pET28a-rpo-gene plasmids were grown in 500 ml of LB medium, supplemented with 25 μg/ml kanamycin, at 37°C until an OD600 of around 1 was reached. Then the culture was induced to express an RNAP subunit by the addition of 1 mM isopropyl β-D-1-thiogalactopyranoside and allowed to grow for 2-4 hours. Cells were harvested by centrifugation and stored at -80°C before use.

Protein purification

Frozen cells were thawed and lysed by sonication in a buffer containing 40 mM Tris-HCl, pH 8.0, 0.1 M NaCl, 10 mM EDTA, 10 mM 2-ME. The lysate was clarified by centrifugation for 30 min at 15,000 × g, and
supernatant was supplemented with ammonium sulphate to precipitate soluble proteins (α2, α1His6, α2His6). Pellets containing insoluble proteins (α1, β, α1His6, α2His6, His6-β′) were resuspended in 40 mM Tris-HCl, pH 8.0, 0.3 M KCl, 10 mM EDTA, 0.2% Nadeoxycholate to prepare inclusion bodies as described before [22,42]. Ammonium sulphate precipitate was collected by centrifugation, and the pellet, containing untagged protein (α2), was dissolved in 40 mM Tris-HCl, pH 8.0, 0.1 M NaCl, 1 mM EDTA, 2 mM 2-ME, 0.1 mM PMSF and loaded onto a 1 ml HiTrap Heparin HP column (GE Healthcare) equilibrated in the same buffer. The bound protein was eluted by a linear gradient of NaCl (from 0.1 to 1.0 M) in the same buffer. The pellet containing α1His6 or α2His6 was dissolved in 20 mM Tris-HCl, pH 8.0, 0.5 M NaCl, 2 mM 2-ME, 0.1 mM PMSF, 5 mM imidazole and loaded onto the 1 ml HiTrap Cheating HP column (GE Healthcare) charged with Ni2+. The column was washed and bound protein was step-eluted with 20, 50, or 100 mM imidazole. Fractions containing pure protein were pooled and dialyzed against two changes of 500 volumes of 40 mM Tris-HCl, pH 8.0, 100 mM NaCl, 0.5 mM EDTA, 0.5 mM 2-ME, 10% glycerol, concentrated by ultrafiltration using Microsep (PALL, Life Science) centrifugal device, and stored at -80°C.

In vitro protein interaction experiments
Purified proteins were mixed together in pairwise combinations of untagged and his-tagged proteins. Before mixing, proteins in inclusion bodies were solubilised in denaturing buffer containing 6M guanidine-HCl, 20 mM Tris-HCl, pH 8.0, 10 mM MgCl2, 10 μM ZnCl2, 1 mM EDTA, 10 mM DTT, 10% glycerol. Coupled proteins were mixed in 0.5-1 ml of denaturing buffer at equimolar ratio and adjusted to 0.2-0.5 mg/ml total protein concentration. Refolding of denatured molecules was achieved by removing guanidine-HCl from reaction mix through one-change dialysis against 500 ml of reconstitution buffer (20 mM Tris-HCl, pH 8.0, 0.2 M NaCl, 10 mM MgCl2, 10 μM ZnCl2, 0.25 mM EDTA, 0.5 mM DTT, 20% glycerol). Precipitate formed during dialysis was removed by centrifugation. Then supernatant was diluted 4-fold with start buffer (25 mM HEPES, pH 8.0, 0.5 M NaCl, 5% glycerol, 1 mM imidazole) and loaded on 0.5 ml His-Select Nickel Affinity Gel (Sigma) column equilibrated in the same buffer. The column was washed with start buffer, and proteins were eluted with three steps of start buffer containing 20, 100, 200 mM imidazole. Fractions were analyzed by SDS-PAGE and visualized by Coomassie-staining.

Reconstitution of α2β, and α2β′ was performed as described above. The α2β subassemblies were stepwise fractionated on Ni2+-affinity column with 10, 50, and 100 mM imidazole; α2β′ RNAP core assembly reactions were fractionated by gel-filtration on a Superose 6 column (GE Healthcare) in the buffer containing 40 mM Tris-HCl, pH 8.0, 100 mM NaCl, 1 mM EDTA, and 1 mM 2-ME. Superose 6 fractions were checked for transcription activity on a nucleic acid scaffold. For this purpose, nucleic acid scaffold containing radioactively labelled 8 nt RNA primer (Figure 3A) was added into 10 μl of the target Superose 6 fraction to obtain artificial transcription elongation complexes and transcription was initiated by the addition of NTP and Mg2+. Reaction products were resolved by denaturing 20% PAGE and revealed by autoradiography.

Bacterial two-hybrid assays
Gene fragments encoding Francisella α1NTD (residues 1-244), α2NTD (residues 1-242), α1CTD (residues 227-323), α2CTD (residues 262-318), were cloned in the plasmids pBRαLN and pACαCl to gain fusions with E. coli RNAP αNTD and bacteriophage λ cl proteins respectively [24,43,44]. FW102 F′O2-26 reporter strain cells were co-transformed with every possible pairwise plasmid combination. Individual transformants were selected and grown in 2 ml of LB medium supplemented with 50 μg/ml carbenicillin, 25 μg/ml kanamycin, 25 μg/ml chloramphenicol, and 0.1 mM IPTG. β-galactosidase assays were performed as described earlier [43].

RpoA comparative analysis
For comparative analysis of two RpoA paralogs in Francisella we retrieved the RefSeq database (NCBI) containing 1055 completely sequenced bacterial genomes on March 2010. A set of 368 RpoA sequences from 355 representative genomes was aligned with MUSCLE program [45], and the maximum likelihood tree for 246 informative aligned positions was built using FastTree program [46]. RAxML program [31] was used for reconstruction of the phylogenetic tree for γ-proteobacteria with β-proteobacteria as an outgroup. The same program was used for comparison of the maximum likelihood values for the best and constrained trees. The evolutionary model for tree reconstruction (WAG [47] with gamma-distributed evolutionary rates) was selected with ProtTest program [48].

Additional material

Additional file 1: Alpha alignment
Additional file 2: Constrained trees

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