Structures of the Noncanonical RNA Ligase RtcB Reveal the Mechanism of Histidine Guanylylation

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Supporting Information

ABSTRACT: RtcB is an atypical RNA ligase that joins either 2′,3′-cyclic phosphate or 3′-phosphate termini to 5′-hydroxyl termini. In contrast to classical RNA ligases, which rely on ATP and Mg(II), catalysis by RtcB is dependent on GTP and Mn(II) with ligation proceeding through a covalent RtcB–histidine–GMP intermediate. Here, we present three structures of Pyrococcus horikoshii RtcB complexes that capture snapshots along the entire guanylylation pathway. These structures show that prior to binding GTP, a single manganese ion (Mn1) is bound to RtcB. To capture the step immediately preceding RtcB guanylylation, we determined a structure of RtcB in complex with Mn(II) and the unreactive GTP analogue guanosine 5′-(α-thio)triphosphate (GTPαS). This structure shows that Mn1 is poised to stabilize the pentavalent transition state of guanylylation while a second manganese ion (Mn2) is coordinated to a nonbridging oxygen of the γ-phosphoryl group. The pyrophosphate leaving group of GTPαS is oriented apically to His404 with the ε-nitrogen poised for in-line attack on the α-phosphorus atom. The structure of RtcB in complex with GTPαS also reveals the network of hydrogen bonds that recognize GTP and illuminates the significant conformational changes that accompany the binding of this cofactor. Finally, a structure of the enzyme histidine–GMP intermediate depicts the end of the guanylylation pathway. The ensuing molecular description of the RtcB guanylylation pathway shows that RtcB and classical ATP- and Mg(II)-dependent nucleic acid ligases have converged upon a similar two-metal mechanism for formation of the nucleotidylated enzyme intermediate.

RNA ligases catalyze the formation of a phosphodiester bond between RNA termini that are generated by specific endonucleases during tRNA splicing, the unfolded protein response, and the antiphage response.1–4 These endonucleases generate 2′,3′-cyclic phosphate and 5′-OH termini upon cleavage.5,6 Classical ATP-dependent RNA ligases in bacteria, fungi, and plants are components of multi-enzyme pathways that repair RNAs with 2′,3′-cyclic phosphate and 5′-OH ends.7,8 Before ligation, the 2′,3′-cyclic phosphate is hydrolyzed to a 3′-OH by a phosphodiesterase and the 5′-OH is phosphorylated by a polynucleotide kinase to generate a 5′-phosphate (5′-P). Classical ligases then catalyze the ATP- and Mg(II)-dependent joining of 5′-P and 3′-OH termini.1

The noncanonical RNA ligase RtcB catalyzes an unprecedented reaction, joining 2′,3′-cyclic phosphate and 5′-OH RNA termini.9–18 RtcB is an essential enzyme for the maturation of tRNAs in metazoa13 and possibly archaea12 and shares no sequence or structural similarity16 with canonical nuclear acid ligases. In marked contrast to classical ligases, RtcB relies on GTP and Mn(II) for catalysis. Ligation proceeds through three nucleotidyl transfer steps, with 2′,3′-cyclic phosphate termini being hydrolyzed to 3′-P termini in a step that precedes 3′-P activation with GMP (Figure 1A).14,16,17 In the first nucleotidyl transfer step, RtcB reacts with GTP to form a covalent RtcB–histidine–GMP intermediate and release PPi; in the second step, the GMP moiety is transferred to the RNA 3′-P, and in the third step, the 5′-OH from the other RNA strand attacks the activated 3′-P to form a phosphodiester bond and release GMP. Thus, a high-energy phosphoanhydride of GTP activates a 3′-P for intermolecular attack by a 5′-OH. Here, we provide insight into the chemical mechanism of an unusual nucleotidyl transfer reaction in this sequence, Mn(II)-dependent histidine guanylylation.

We sought to elucidate the entire pathway of RtcB guanylylation by determining the three-dimensional structures of key intermediates at atomic resolution. We present three structures of Pyrococcus horikoshii RtcB complexes. (i) A structure with bound Mn(II) represents the intermediate that precedes binding of GTP. (ii) A structure with bound Mn(II) and an unreactive GTP analogue, guanosine 5′-(α-thio)triphosphate (GTPαS), captures the reaction step immediately preceding formation of the covalent enzyme intermediate. (iii) A structure of the covalent RtcB–histidine–GMP intermediate depicts the end product of the guanylylation pathway.
results show that RtcB coordinates a single Mn(II) ion prior to binding GTP and that GTP binds to RtcB in a complex with a second Mn(II) ion. This two-manganese mechanism of RtcB guanylylation is analogous to the two-magnesium mechanism of adenylylation used by canonical ATP-dependent nucleic acid ligases.20,21

■ MATERIALS AND METHODS

RtcB Purification. A previously described plasmid expressing P. horikoshii RtcB was used except that the sequence encoding the hexahistidine tag was removed via mutagenesis.15 Native P. horikoshii RtcB was expressed in BL21 cells when they were grown in Terrific Broth at 37 °C to an OD_{600} of 0.6, induced with IPTG (0.5 mM), and grown for an additional 3 h. Cells were harvested by centrifugation and resuspended at 8 mL per gram of wet pellet in buffer A [50 mM MES-NaOH (pH 5.6), 45 mM NaCl, and 1 mM Cleland’s reagent]. Cells were lysed by being passage through a cell disruptor (Constant Systems) at 20000 psi, and the lysate was clarified by centrifugation at 20000 g for 25 min followed by centrifugation at 20000 g for 20 min. The clarified lysate was then loaded onto a 5 mL HiTrap HP SP cation-exchange column (GE Lifesciences). The column was washed with 25 mL of buffer A, and RtcB was eluted with a NaCl gradient of buffer A (45 mM to 1.0 M) over 20 column volumes. Fractions containing RtcB were dialyzed against 4 L of buffer [10 mM HEPES-NaOH (pH 7.5) and 200 mM NaCl] overnight at 4 °C.

14C-Labeled GTP Binding Assays. Binding assays were performed in 250 μL of 50 mM HEPES buffer (pH 7.5) containing NaCl (200 mM), P. horikoshii RtcB (100 μM), various concentrations of MnCl_{2}, and [8-14C]GTP (Moravek Biochemicals, Brea, CA).22 After incubation, free GTP was removed by applying the reaction mixture to three 5 mL HiTrap desalting columns (GE Lifesciences) connected in series. The desalting columns were equilibrated with elution buffer [50 mM HEPES (pH 7.5) and 200 mM NaCl], and protein was eluted in 0.5 mL fractions. Absorbance readings at 2519 and 2525 readings for each fraction. The protein fractions have high A_{2519} readings, whereas the fractions with free GTP have higher A_{2525} readings. In fractions containing

Figure 1. RtcB stepwise ligation reaction pathway and electron density maps of the RtcB active sites. (A) RtcB joins RNA in three nucleotidyl transfer steps, with 2′,3′-cyclic phosphate termini being hydrolyzed in a step preceding RNA 3′-P activation. The three nucleotidyl transfer steps are (1) RtcB guanylylation, (2) RNA 3′-P activation, and (3) phosphodiester bond formation. The RNA 3′-P at the ligation junction is colored red to ease visualization of its incorporation into the newly generated 3′,5′-phosphodiester bond. (B) RtcB−Mn(II) complex with red mesh representing a composite simulated annealing omit map contoured at 5.6σ to show the presence of one Mn(II) ion. (C) RtcB−GTPo5−Mn(II) complex with blue mesh denoting an omit map (F_o−F_c) of the GTPo5 density contoured at 2.6σ. (D) RtcB−GMP−Mn(II) complex with blue mesh denoting an omit map (F_o−F_c) of the GMP density contoured at 3.5σ.
protein, the RtcB concentrations were calculated from the $A_{280}$ reading using an extinction coefficient of 62340 M$^{-1}$ cm$^{-1}$ (ExPASy). The concentration of $[8-14C]$GTP in the protein fractions was determined by liquid scintillation counting. Each 0.5 mL fraction was mixed with 3.5 mL of Ultima Gold MV liquid scintillation cocktail (Perkin-Elmer) in a 4 mL vial, and counts were read on a MicroBeta TriLux liquid scintillation counter (Perkin-Elmer). The concentration of GTP in each fraction was determined by comparing the counts per minute in these samples to the values obtained from standards with known concentrations. Optimal formation of the RtcB–GMP complex was found to occur in reaction mixtures that included 1 mM GTP and 2 mM MnCl$_2$. The optimal incubation conditions were found to be at 70 °C for 45 min. Under these conditions, the GTP:RtcB molar ratio was determined to be (0.76 ± 0.02):1. No binding of GTP to RtcB was detected in the absence of Mn(II).

Figure 2. Wall-eyed stereoviews of the active sites of RtcB complexes (subunit A). (A) RtcB–Mn(II) complex [Protein Data Bank (PDB) entry 4isj]. (B) RtcB–GTPαS–Mn(II) complex (PDB entry 4isz). GTPαS is modeled as the R$_p$ diastereomer. (C) RtcB–GMP–Mn(II) covalent intermediate (PDB entry 4it0). Hydrogen bonding to the guanosine ribose 2’-OH and 3’-OH with the backbone amides of Ala406 and Gly407, respectively, is not shown.
Accelerated Publication

**RtcB Crystallization.** RtcB was concentrated to 200 μM (11 mg/mL) by ultrafiltration using a spin concentrator (5000 molecular weight cutoff, Amicon) and passed through a 0.2 μm filter. To prepare the RtcB−Mn(II) complex, MnCl₂ (1 mM) was added to the concentrated protein. For preparation of the RtcB−GTPαS−Mn(II) complex, MnCl₂ (2 mM) and a 1:1 mixture of Rₐ and Sₐ diastereomers of GTPαS (1 mM) was added to the concentrated protein solution, and the resulting solution was incubated at 70 °C for 15 min. For preparation of the RtcB−GMP−Mn(II) complex, the covalent intermediate was formed as described above, and the solution was subjected to gel-filtration chromatography on a Superdex 16/60 column (GE Lifesciences) to remove P₈ and excess MnCl₂ and GTP. Each of the protein complexes was flash-frozen in liquid nitrogen and stored at −80 °C. Protein samples were crystallized using the hanging drop vapor diffusion method. Crystals were grown by mixing 1 μL of sample solution with 1 μL of reservoir solution. The RtcB−Mn(II) and RtcB−GTPαS−Mn(II) complexes were crystallized using identical reservoir solutions consisting of Bis-Tris (0.1 M, pH 5.5) and ammonium sulfate (2.1 M) and the RtcB−GMP−Mn(II) complex was crystallized using HEPES-NaOH (0.1 M, pH 7) and ammonium sulfate (2 M). Trays were incubated at 20 °C, and crystals appeared within 1 week. Crystals were harvested and cryoprotected in a reservoir solution containing sucrose [20% (w/v)] and cryopreserved in liquid nitrogen.

**Data Collection, Structure Determination, and Refinement.** X-ray diffraction data were collected at 100 K at the Life Science Collaborative Access Team at the Advanced Photon Source at Argonne National Laboratory (Argonne, IL). Data sets were indexed and scaled using HKL2000. The apo-RtcB structure was used as a starting model, and the structures were completed using alternating rounds of manual model building using COOT and refinement with phenix.refine. Structure quality was assessed by MolProbity, and structure factors were calculated using Phenix.

## RESULTS

**A Structure with Mn(II) Represents the Intermediate That Precedes GTP Binding.** For crystallization of the RtcB−Mn(II) complex, MnCl₂ (1 mM) was added to the concentrated protein solution (200 μM) before crystallization. That Precedes GTP Binding. For crystallization of the RtcB−GTPαS−Mn(II) complex, MnCl₂ (2 mM) and GTPαS (1 mM) were added to a concentrated solution of protein (200 μM), and the resulting solution was incubated at 70 °C for 15 min to facilitate any conformational changes in the hyperthermophilic enzyme that are necessary for cofactor binding. The RtcB−GTPαS−Mn(II) complex was subsequently maintained at temperatures that disallow formation of the covalent intermediate. Crystals of this complex diffracted to an effective resolution of 2.45 Å (1/σ₁ = 2); however, all data to 2.3 Å were used in refinement (Table S1 of the Supporting Information).

The omit density map of the RtcB−GTPαS−Mn(II) complex indicated the presence of GTPαS and two manganese ions in the RtcB active site (Figure S1B of the Supporting Information and Figures 1C and 2B). Mn1 remains in a tetrahedral coordination geometry with ligands that include the same three amino acid residues as the structure with manganese only; however, the water molecule has been replaced with the nonbridging α-thiophosphate oxygen of GTPαS. A second manganese ion (Mn2) is in a tetrahedral coordination geometry with ligands that include a nonbridging oxygen of the γ-phosphoryl group of GTPαS, as well as three amino acid residues (Asp95, Cys98, and His203). The β-phosphoryl group of GTPαS is oriented apically to His404, and its N⁰ is poised for in-line attack on the α-phosphoryl group. Furthermore, H-N⁰ of His404 forms a hydrogen bond with O⁹¹ of Asp65, which is strictly conserved and appears to orient and activate N⁰ for attack. A main chain H-N forms a hydrogen bond with O⁶² of Asp65, stabilizing an anti orientation of the carboxylate in the His--Asp dyad.

The presence of a cysteine residue bridging two manganese ions in the RtcB active site is unique. The Mn1...S and Mn2...S coordination distances in the RtcB−GTPαS−Mn(II) complex are 2.3 and 2.4 Å, respectively, and the Mn1...S−Mn2 angle is 100°. Thus, the two Mn(II) ions are separated by only 3.6 Å. This distance is similar to the 3.3 Å distance separating the Mn(II) ions of the renowned binuclear manganese cluster in the active site of arginase, where the Mn(II) ions are bridged by two aspartate residues. The manganese ion coordination distances in the RtcB−GTPαS−Mn(II) complex are listed in Table 1.

RtcB is the only known enzyme catalyzing nucleotidyl transfer that requires a NTP−Mn(II) complex rather than a NTP−Mg(II) complex as a cofactor. Indeed, RtcB is not active with Mg(II), which is much more abundant than Mn(II) in both cells and the environment. The structure of the RtcB−GTPαS−Mn(II) complex provides an explanation for this unusual requirement. First, the ligands of the two bound Mn(II) ions have a tetrahedral geometry, which is disfavored by Mg(II). Second, the side chain of a cysteine residue interacts with both Mn(II) ions, which are more thiophilic than Mg(II) ions. This essential cysteine residue is strictly conserved

dx.doi.org/10.1021/bi4002375 | Biochemistry 2013, 52, 2518–2525

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throughout evolution and likely serves as a gatekeeper that selects for Mn(II) in each metal-binding site. Third, Coulombic repulsion deters the close placement of two Mg(II) ions, which have a high charge density. Indeed, the two Mg(II) ions used by T4 RNA ligase are separated by 7.4 Å, a distance that is 2-fold greater than that of the Mn(II) ions in RtcB. The two Mg(II) ions in the active site of xylose isomerase, which are bridged by a glutamate carboxylate, have a shorter internuclear distance of 5.1 Å. More polarizable Mn(II) ions, however, can be accommodated in even greater proximity.

An intricate array of hydrogen bonds explains the specificity and high affinity for GTP. The triphosphate moiety forms hydrogen bonds with H-Nδ of two asparagine residues. Asn202 has now adopted a different conformation, and its H-Nδ forms a hydrogen bond with the β-phosphoryl group. Likewise, the H-Nδ of Asn330 forms a hydrogen bond with the γ-phosphoryl group. The guanosine nucleoside is bound in an anti conformation with the guanine base stacked on Phe204 and with Tyr451 forming an edge of the guanine-binding pocket. Each carboxylate oxygen of Glu206 forms a hydrogen bond with guanine, one with the H-N1 and the other with the H-N2;
Ser385 also interacts with the H-N2, while the H-N of Lys480 forms a hydrogen bond with O6. The guanosine ribose 2'- and 3'-oxygen atoms form hydrogen bonds with the main chain H-N of Ala406 and Gly407, respectively.

The binding of GTPαS elicits significant conformational changes in the RtcB active site (Figure 3A). The loop that is displaced by the guanine base has a maximal Cα displacement of 2.5 Å at Ser380. In addition, the loop containing Ala406 and Gly407 changes conformation around the ribose 2'-OH and 3'-OH with a maximal Cα displacement of 1.4 Å at Ala406.

**Structure of the RtcB—Histidine—GMP Covalent Intermediate.** To determine the optimal reaction conditions that allow formation of the RtcB—GMP covalent intermediate, 14C-labeled GTP binding studies were performed. The optimal reaction conditions were found to include purified RtcB (100 μM), GTP (1 mM), and MnCl₂ (2 mM), with incubation at 70 °C for 45 min. Under these conditions, the maximal GMP-RtcB molar ratio was determined to be (0.76 ± 0.02):1. No binding of GTP to RtcB was detected in the absence of GTP, and PPi by gel-maximal GMP:RtcB molar ratio was determined to be (0.76 ± 0.02):1. No binding of GTP to RtcB was detected in the absence of Mn(II). Using these reaction conditions, we formed the RtcB—GMP intermediate and removed unbound Mn(II), GTP, and PPi, by gel-filtration chromatography. The protein was concentrated to 200 μM, and crystals of this complex diffracted to a resolution of 2.4 Å (Table S1 of the Supporting Information).

The omit density map indicated the presence of a covalent histidine—GMP complex and two manganese ions in the RtcB subunit A active site (Figure S1C of the Supporting Information and Figures 1D and 2C). The GMP density present in subunit B was too weak to model confidently. The guanine and ribose interactions are essentially identical to those in the RtcB—GTPαS—Mn(II) complex. Asn202, however, has shifted to a position near the nascent phosphoramidate bond (Figure 3B). The labile histidine—GMP complex is stabilized by coordination of one nonbridging oxygen of the GMP to Mn1 and the formation of a hydrogen bond of a hydroxyl oxygen of the other nonbridging oxygen with a water molecule. The phosphoimidazolium group of His404 is stabilized by a hydrogen bond from its H-N of the carboxylate side chain of Asp65. Mn2 remains bound in a tetrahedral coordination geometry; however, the metal contact with the γ-phosphoryl group has been replaced with a water molecule. Alanine-scanning mutagenesis of GMP-interacting residues revealed their importance for RNA ligation activity, consistent with a previous report (Figures S2 and S3 of the Supporting Information).

**Discussion**

**RtcB Guanylylation Mechanism.** Histidine guanylylation is expected to proceed through an associative mechanism with the accumulation of negative charge on the nonbridging oxygens of the α-phosphoryl group in the pentavalent transition state. In the RtcB active site, guanylylation is promoted by neutralization of this negative charge by coordination to Mn1 and hydrogen bonds with water molecules. The PPi leaving group apical to Nδ of His404 on the α-phosphorus of GTP is promoted by (i) orientation and stabilization of His404 by a hydrogen bond with the H-N of Asp65, (ii) orientation of the PPi, leaving group apical to Nδ of His404 by Mn2, and (iii) charge neutralization of the pentavalent transition state by Mn1 and two water molecules. Two-metal mechanism of adenylylation used by T4 RNA ligase based on a structure in complex with magnesium ions and the nonhydrolyzable ATP analogue ApCpp (PDB entry 2c5u). Hydrogen bonds with ApCpp are not shown, and only the inner sphere ligands of Mg1 and Mg2 are shown. In the crystal structure, the Mg1 site was occupied by a Ca(II) ion, which was present at a high concentration in the crystallization solution. Mg1 and Mg2 serve mechanistic roles essentially identical to those of Mn1 and Mn2, respectively, in the RtcB guanylylation mechanism.

A structure of T4 RNA ligase bound to the ATP analogue adenosine 5′-(α,β-methylene)triphosphate (ApCpp) is consistent with a mechanism analogous to the one described here for RtcB guanylylation. In the T4 RNA ligase structure, a calcium ion is bound in place of one magnesium ion (Mg1) and coordinates to a nonbridging oxygen of the ApCpp α-phosphate and a magnesium ion (Mg2) coordinates to the β-phosphonate group. Mg1 in the T4 ligase structure and Mn1 in the RtcB structure both promote enzyme nucleotidylation by neutralizing the negative charge on the α-phosphoryl group in the pentavalent transition state. The second metal ion observed
in both T4 ligase and RtcB enters the active site in a coordination complex with the NTP cofactor and promotes catalysis by orienting the PP, leaving group and neutralizing the charge on the phosphoryl groups. Despite the absence of sequence or structural similarity between RtcB and classical ATP-dependent nucleic acid ligases, Nature has converged on analogous two-metal-dependent nucleotidylation mechanisms.

The structures presented here resolve key issues about RtcB guanylylation. In particular, the structure of the RtcB−GTPαS−Mn(II) complex has revealed the orientation of the bound triphosphate cofactor and the role of Mn2. The consequent molecular description of the two-metal RtcB guanylylation mechanism contrasts with one proposed previously,18 which depicted an incorrect orientation for its β- and γ-phosphoryl groups.

All known nucleotidyl transferases require complexation of a nucleotide triphosphate cofactor to a metal ion. The roles of the metal ion in the NTP−metal complex include orientating the phosphoryl groups, neutralizing their charge, and enhancing their reactivity.36,37 Many enzymes that catalyze the cleavage of an NTP α−β phosphoanhydride bond employ two Mg(II) ions, one that coordinates to the high-affinity site between the β- and γ-phosphoryl groups and a second that coordinates to the α-phosphoryl group.38−43 We have revealed how an enzyme can, instead, employ two Mn(II) ions in analogous roles to catalyze nucleotidyl transfer.

■ ASSOCIATED CONTENT

* Supporting Information

Active site electron density (2Fo−Fc) of refined models (Figure S1), structure-guided mutagenesis of the guanylate-binding pocket (Figures S2 and S3), and crystallographic data collection and refinement statistics (Table S1). This material is available free of charge via the Internet at http://pubs.acs.org.

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Funding

This work was supported in part by National Institutes of Health Grants F32 GM100681 (to K.K.D.); Protein Structure Initiative Grants U01 GM098248 (to G.N.P.) and U54 GM074901 (Center for Eukaryotic Structural Genomics); and R01 CA073808 (to R.T.R.). The Life Sciences Collaborative Access Team has been supported by Michigan Economic Development Corp. and the Michigan Technology Tri-Corridor. Use of the Advanced Photon Source was supported by the U.S. Department of Energy, Basic Energy Sciences, Office of Science, under Contract W-31-102-ENG-38.

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

We are grateful to Prof. Aaron A. Hoskins for helpful discussions.

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