Novel Mechanism of RNA Repair by RtcB via Sequential 2',3'-Cyclic Phosphodiesterase and 3'-Phosphate/5'-Hydroxyl Ligation Reactions*5

Received for publication, September 8, 2011, and in revised form, October 27, 2011 Published, JBC Papers in Press, October 31, 2011, DOI 10.1074/jbc.M111.302133

Naoko Tanaka, Anupam K. Chakravarty, Bill Maughan, and Stewart Shuman1

From the Molecular Biology Program, Sloan-Kettering Institute, New York, New York 10065

Background: RtcB is a new RNA ligase implicated in tRNA splicing and RNA repair.

Results: RtcB has 2',3'-cyclic phosphodiesterase and 3'-phosphate/5'-hydroxyl ligation activities.

Conclusion: RtcB seals broken RNAs via a two-step reaction pathway.

Significance: The structure, active site, and chemical mechanism of RtcB are unique.

RtcB enzymes are a newly discovered family of RNA ligases, implicated in tRNA splicing and other RNA repair reactions, that seal broken RNAs with 2',3'-cyclic phosphate and 5'-OH ends. Parsimony and energetics would suggest a one-step mechanism for RtcB sealing via attack by the O5' nucleophile on the cyclic phosphate, with expulsion of the ribose O2' and generation of a 3',5'-phosphodiester at the splice junction. Yet we find that RtcB violates Occam’s razor, insofar as (i) it is adept at ligating 3'-monophosphate and 5'-OH ends; (ii) it has an intrinsic 2',3'-cyclic phosphodiesterase activity. The 2',3'-cyclic phosphodiesterase and ligation reactions both require manganese and are abolished by mutation of the RtcB active site. Thus, RtcB executes a unique two-step pathway of strand joining whereby the 2',3'-cyclic phosphodiester end is hydrolyzed to a 3'-monophosphate, which is then linked to the 5'-OH end to form the splice junction. The energy for the 3'-phosphate ligation activity is provided by GTP, which reacts with RtcB in the presence of manganese to form a covalent RtcB-guanylate adduct. This adduct is sensitive to acid and hydroxylamine but resistant to alkali, consistent with a phosphoramidate bond.

RNA 2',3'-cyclic phosphate and 5'-OH ends figure prominently in RNA metabolism: as the products of RNA cleavage by site-specific endoribonucleases, as substrates for enzymatic RNA end joining during tRNA repair and tRNA splicing, and as substrates for unconventional mRNA splicing during the unfolded protein response (1–6). Broadly speaking, there are two known RNA repair pathways that rectify 2',3'-cyclic phosphate/5'-OH breaks. The “healing-and-sealing” pathway of RNA repair is a multistep process typically catalyzed by three different enzymes dedicated to 3’ healing, 5’ healing, and ligation, respectively (6, 7). In the healing phase, the 2',3'-cyclic phosphate end is hydrolyzed to a 3'-OH, and the 5'-OH end is phosphorylated by an NTP-dependent polynucleotide kinase to yield a 5'-monophosphate. The healed 3’-OH and 5’-PO4 termini are then suitable substrates for a “classic” ATP-dependent RNA ligase that restores the 3’, 5’ phosphodiester backbone via three nucleotidyl transfer steps as follows: (i) ligase reacts with ATP to form a ligase-(lysyl-N)-AMP intermediate plus pyrophosphate, (ii) AMP is transferred from ligase-adenylate to the 5’-PO4 RNA end to form an RNA-adenylate intermediate, AppRNA,2 and (iii) ligase catalyzes attack by an RNA 3’-OH on AppRNA to form a 3’, 5’ phosphodiester bond and release AMP (8). The salient principle of classic ligase catalysis is that the high energy of the ATP phosphoanhydride bond is transferred via the enzyme to the RNA 5’ end, thereby activating the 5’ end for phosphodiester synthesis.

RNA end-healing enzymes and classic RNA ligases are found in diverse taxa in all three phylogenetic domains. The healing-and-sealing pathway is responsible for tRNA splicing in fungi and plants (9, 10), for mRNA splicing in the fungal unfolded protein response (4, 5), and for tRNA restriction-repair during bacteriophage infection of Escherichia coli (3). Recent studies highlight a novel bacterial healing-and-sealing pathway that is thought to confer adaptive immunity to tRNA damage inflicted by ribotoxins (11, 12).

A second pathway of RNA repair by “direct ligation” of 2',3'-cyclic phosphate and 5'-OH ends was discovered nearly 30 years ago via analysis of the mammalian tRNA splicing reaction (13, 14). Specifically, it was shown that the 2',3'-cyclic phosphate of the cleaved pre-tRNA is incorporated at the splice junction in the mature tRNA. (By contrast, the junction phosphate in yeast tRNA splicing derives from the γ-phosphate of the NTP substrate for the 5’ kinase step.) Direct ligation is thought to be the main pathway of tRNA splicing in animals and archaea (13–15). Earlier this year, three laboratories reported the identification of bacterial, archaeal, and mammalian RtcB proteins as novel RNA ligase enzymes capable of seal ing 2',3'-cyclic phosphate and 5'-OH ends in the absence of ATP (16–18). RtcB homologs are present in metazoan and prok oza but not in fungi and plants. Thus, the phylogenetic distri-

---

* This work was supported, in whole or in part, by National Institutes of Health Grant GM46330.

5 The on-line version of this article (available at http://www.jbc.org) contains supplemental Figs. S1–S3.

1 An American Cancer Society Research Professor. To whom correspondence should be addressed. E-mail: s-shuman@ski.mskcc.org.

2 The abbreviations used are: AppRNA, RNA-adenylate intermediate; Pnkp, T4 polynucleotide kinase-phosphatase; AtRNL, plant tRNA ligase; CPDase, 2',3'-cyclic phosphodiesterase; ChV-Lig, Chlorella virus DNA ligase; 2',3’ > p, 2',3’ cyclic phosphate.
Two-step Pathway of End-joining by RNA Ligase RtcB

One-step direct ligation of 2',3'-p and 5'-OH ends
Two-step pathway of cyclic phosphodiester hydrolysis and 3'-P/5'-OH ligation

FIGURE 1. Alternative models for RtcB catalysis. A, one-step direct ligation; B, two-step pathway.

The role of RtcB in RNA repair is of broad interest, as it is a key enzyme in the repair of broken RNA molecules.

**Experimental Procedures**

**Purification of RtcB**—pET28b-His10Smt3-RtcB plasmids encoding wild-type RtcB and mutants C78A and H168A were transformed into *E. coli* BL21-CodonPlus(DE3). Cultures (1 liter) derived from single kanamycin-resistant transformants were grown at 37 °C until the A_{590} reached 0.6–0.8. The cultures were chilled on ice for 30 min and then adjusted to 0.1 mM isopropyl β-D-thiogalactoside. Incubation was continued at 17 °C for 16 h with constant shaking. Cells were harvested by centrifugation and stored at −80 °C. Preparation of soluble lysates, recovery of the His_{10}Smt3-RtcB proteins by nickel-agarose chromatography, excision of the tags by treatment with the Smt3-specific protease Ulp1, separation of the tag-free RtcB proteins from His_{10}Smt3 by a second round of Ni-agarose chromatography, and final purification of RtcB to by Superdex 200 gel filtration were performed as described previously (18). Protein concentrations were determined using the Bio-Rad dye reagent with BSA as the standard.

**RNA Repair Substrates**—3'-Phosphorylated RNA oligonucleotides labeled with ^32P were prepared by T4 RNA ligase 1 (Rnl1)-mediated addition of [5',32P]pCp to a 19-mer synthetic oligoribonucleotide. The [5',32P]pCp donor was generated via enzymatic phosphorylation of unlabeled 3'-CMP by T4 polynucleotide kinase-phosphatase (Pnkp) mutant D167N (kinase active, but devoid of 3'-phosphatase; purified according to Ref. 22) in the presence of [γ-32P]ATP. A kinase reaction mixture (20 μl) containing 50 mM Tris-HCl (pH 8.0), 10 mM MgCl_2, 10 mM DTT, 0.1 mM 3'-CMP, 0.1 mM [γ-32P]ATP (130 μCi), 0.1 mg/ml BSA, and 10 μg/ml Pnkp-D167N was incubated at 37 °C for 60 min. The Pnkp was inactivated by heating the mixture at 95 °C for 3 min. The mixture was then adjusted to 40 μl and, in the process, supplemented with 100 μM fresh unlabeled ATP, 50 μM 19-mer oligoribonucleotide 5'-rUGGCUCCGAUAAUC-ACGCUU, and 0.2 mg/ml Rnl1 (purified as described in Ref. 23). The T4 ligase reaction mixture was incubated for 1 h at 37 °C, and the products were resolved by electrophoresis.
Two-step Pathway of End-joining by RNA Ligase RtcB

through a 40-cm 20% polyacrylamide gel containing 8 M urea. The \( ^{32} \text{P} \)-labeled 20-mer oligonucleotide product (5'-\text{HO}UGGCUCCGUAUACCGCUUpCp) was located by autoradiography and excised from the gel. The oligonucleotide was eluted by soaking the gel slice overnight in 0.4 ml of 1 M ammonium acetate, 0.2% SDS, 20 mM EDTA. The eluted oligonucleotide was recovered by ethanol precipitation and resuspended in 10 mM Tris-HCl (pH 6.8).

The purified 20-mer was further modified by 5'-phosphorylation and/or cyclization of the 3'-phosphate as follows. Reaction mixtures (100 \( \mu \)l) containing 50 mM Tris-HCl (pH 7.4), 0.1 mM ATP, 10 mM MgCl2, 2 mM DTT, 0.1 mM spermidine, 500 mM \( ^{32} \text{P} \)-labeled 20-mer RNA, and 10 \( \mu \)g/ml Pnkp-D167N and/or 40 \( \mu \)g/ml E. coli cyclase RtcA (purified according to Ref. 24) were incubated for 30 min at 37 °C. The end-modified reaction products 5'-pUGGCUCCGUAUACCGCUUpCp, 5'-pUGGCUCCGUAUACCGCUUpCp and \( ^{32} \text{P} \)-monophosphate-terminated 20-mer RNA to form the broken stem-loop substrates as specified (formed by mixing 20 nM labeled 2'-P-20-mer with 200 nM unlabeled complementary 20-mer to yield the substrates illustrated at the bottom), and 200 mM RtcB or AtRNL (where indicated by + above the lanes) were incubated for 30 min at 37 °C. The products were resolved by denaturing PAGE and visualized by autoradiography. The radiolabeled RNAs corresponding to the 20-mer substrate strand and the ligated hairpin stem-loop are indicated on the left.

**RESULTS**

RtcB Seals Stem-loops with 3'-Monophosphate/5'-OH Breaks—We showed previously that RtcB was adept at ligating broken stem-loop structures with 2',3'-cyclic phosphate and 5'-OH termini to restore a 3',5'-phosphodiester at the repair junction (18). Here we queried whether RtcB could seal a 3'-monophosphate end. We used T4 RNA ligase 1 to transfer [5'-\( ^{32} \text{P} \)]pCp to the 3' end of a 19-mer RNA oligonucleotide to generate a 20-mer strand (\( ^{32} \text{P} \)-labeled 20-mer RNA oligonucleotide with 5'-OH and 3'-monophosphate ends and a single radiolabel between the 3'-terminal and penultimate nucleotides. We used purified E. coli RNA 3'-phosphate cyclase RtcA (24, 25) to convert the 3'-phosphate end of the labeled 20-mer RNA into a 2',3'-cyclic phosphodiester (\( ^{32} \text{P} \)-labeled 20-mer RNA to form the broken stem-loop substrates as specified (formed by mixing 20 nM labeled 2'-P-20-mer with 200 nM unlabeled complementary 20-mer to yield the substrates illustrated at the bottom), and 200 mM RtcB or AtRNL (where indicated by + above the lanes) were incubated for 30 min at 37 °C. The products were resolved by denaturing PAGE and visualized by autoradiography. The radiolabeled RNAs corresponding to the 20-mer substrate strand and the ligated hairpin stem-loop are indicated on the left.

**Intramolecular Joining of 3'-Monophosphate and 5'-OH Ends**—The radiolabeled 20-mer RNAs were reacted with RtcB in the absence of a complementary strand and the products were analyzed by denaturing PAGE (Fig. 3, top panel). The substrates with 5'-OH ends were efficiently converted to a more
Two-step Pathway of End-joining by RNA Ligase RtcB

FIGURE 3. RtcB catalyzes intramolecular 3' phosphate/5'-OH ligation and has a 2',3'-cyclic phosphodiesterase activity. Reaction mixtures (20 μl) containing 50 mM Tris-HCl (pH 8.0), 0.1 mM GTP, 2 mM MnCl₂, 20 nM 32P-labeled 20-mer RNA strand with 5'-OH, 3' phosphate (3'-P), 3'-phosphate (3'-P), or 2',3'-cyclic phosphate (>P) termini (depicted at the right, with the radiolabeled phosphate denoted by ○), and 200 nM RtcB (where indicated by + above the lanes) were incubated for 30 min at 37 °C. The reactions were quenched by adding EDTA to 4 mM final concentration. Half of the reaction mixture was analyzed directly by denaturing PAGE; the labeled RNAs were visualized by autoradiography (top panel). The positions of the linear substrates, the ligated circles, and multimeric ligation products are indicated on the left. The other half of the reaction mixture was digested with RNase T1. (The relevant sites of RNase T1 incision of the 20-mer substrates and flanking the ligation junction are indicated by arrows above the RNA sequences at the right.) The T1 digests were then resolved by denaturing PAGE and visualized by autoradiography (bottom panel). The positions of the T1 fragments are indicated on the left.

FIGURE 4. RtcB cannot seal 2'-monophosphate/5'-OH RNA breaks. RtcB reaction mixtures (20 μl) contained 50 mM Tris-HCl (pH 8.0), 0.1 mM GTP, 2 mM MnCl₂, 20 nM 32P-labeled 20-mer RNA strands with 5'-OH and either 2',3'-cyclic phosphate (>P) or 2'-monophosphate (2'-P) termini (depicted at the right, with the radiolabeled phosphate denoted by ○), and 200 nM RtcB. AtRNL reaction mixtures (20 μl) contained 50 mM Tris-HCl (pH 8.0), 0.1 mM ATP, 0.1 mM GTP, 2 mM DTT, 10 mM MgCl₂, 20 nM 32P-labeled 20-mer RNA strands, and 200 nM AtRNL. The mixtures were incubated for 30 min at 37 °C, and the reactions were quenched with EDTA. The products were analyzed directly by denaturing PAGE; the labeled RNAs were visualized by autoradiography. The positions of the linear substrates, the ligated circles, and multimeric ligation products are indicated on the left.

rapidly migrating circular RNA species as a consequence of intramolecular ligation (lanes 2 and 4). Low levels of more slowly migrating multimers (products of intermolecular ligation) were also generated. There was little difference in the extents of circularization of 5'-OH terminated single strands with 3'-monophosphate versus cyclic phosphate ends. The two substrates with 5'-PO₄ ends were not sealed by RtcB (lanes 6 and 8).

The input 3' end structures and the RNA splice junctions were characterized further by digesting the samples with RNase T1 prior to their electrophoresis through a polyacrylamide gel (Fig. 3, bottom panel). RNase T1 incised the 20-mer RNA substrates 3' of the most distal guanosine to yield 32P-labeled tetranucleotides: either HOCUUpCp or HOCUUpCp-p - with distinct electrophoretic mobilities (lanes 1 and 3). (This analysis verified that the efficiency of cyclization of the CpUpCp end of the 20-mer during the preparative reaction with RtcA was 95%, i.e. 5% of the RNase T1 product was HOCUUpCp in lane 3.) The key finding was that the reactions with RtcB resulted in depletion of the T1 fragments derived from the substrate strands and the appearance of a new, more slowly migrating T1 fragment (Fig. 3, bottom panel, lanes 2 and 4) that corresponded to a 6-mer oligonucleotide, HOCUUpCpUGp, released by T1 incision at the guanosines flanking the predicted ligation junction (Fig. 3). Note that the T1 junction fragment did not accumulate when RtcB was reacted with non-ligatable substrates with 5'-PO₄ ends (lanes 6 and 8).

We further queried the specificity of RtcB by asking whether the enzyme could join a 2'-monophosphate end to a 5'-OH. The 2'-monophosphate 20-mer substrate was prepared by treating the radiolabeled 5'HGGCUCCGAUAUCACGCUUpCp-strand with the isolated C-terminal end-healing domain of plant tRNA ligase (6), which hydrolyzes the 2',3'-cyclic phosphodiester to yield a 2'-phosphate, 3'-OH terminus. (By digesting the labeled 20-mer with RNase T1 before and after treatment with the plant ligase healing domain, and then analyzing the the T1 fragments by PAGE, we verified that the initial 2',3'-cyclic phosphate terminus had been completely converted to a more rapidly migrating phosphomonoester derivative; not shown.) The radiolabeled 2',3'>p and 2'>phosphate 20-mer strands were then reacted in parallel with RtcB and AtRNL and the products were analyzed by PAGE (Fig. 4). As expected, AtRNL catalyzed efficient sealing of both substrates, yielding a mixture of circular and multimeric ligation products (Fig. 4, lanes 3 and 6). By contrast, RtcB sealed the 2',3'>p strand to yield a predominant circular ligation product (lane 2) but was
Two-step Pathway of End-joining by RNA Ligase RtcB

FIGURE 5. Time course of ligation and CPDase activities. Reaction mixtures (80 μl) containing 50 mM Tris-HCl (pH 8.0), 0.1 mM GTP, 2 mM MnCl2, 20 nm 32P-labeled 20-mer RNAs as shown (pRNA in panel A; pRNAp in panel B; pRNA in panel C; pRNAp in panel D), and 200 nm RtcB were incubated at 37 °C. Aliquots (10 μl) were withdrawn at the indicated times (minutes) and quenched immediately with EDTA. The time 0 sample was taken prior to adding RtcB. The quenched mixtures were digested with RNase T1 and then analyzed by denaturing PAGE. The positions of the 3′-P T1 fragment, the 2′,3′-p T1 fragment, and the ligation junction T1 fragment are indicated on the right.

unreactive with the 2′-monophosphate substrate (lane 5). The slight differences in electrophoretic mobility of the circular products made by AtRNL versus RtcB reflect the distinct outcomes of the RtcB and AtRNL ligation reactions. Whereas RtcB seals the 2′,3′-p end to yield a splice junction with a conventional 3′→5′ phosphodiester linkage (18), AtRNL sealing at either a 2′,3′-p or 2′-phosphate end yields a splice junction that retains the 2′-phosphate (1). We conclude that RtcB and AtRNL have antipodal RNA end requirements for sealing, indicative of stark differences in the chemistry of end joining.

RtcB Has 2′,3′-Cyclic Phosphodiesterase Activity—The RNase T1 analysis provided the initially surprising observation that incubation of RtcB with the 5′-phosphorylated 20-mer strand with a cyclic phosphate end, although unproductive with respect to ligation, resulted in a chemical transformation at the 3′ terminus, whereby the original 2′,3′-p was converted to a 3′-monophosphate (Fig. 3, bottom panel, lanes 7 and 8). Thus, in the absence of a polynucleotide 5′-OH, RtcB catalyzes hydrolytic attack on the cyclic phosphodiester. In this experiment, 75% of the input cyclic phosphate was hydrolyzed. We also observed that incubation of RtcB with the 5′-phosphorylated 20-mer strand with a 3′-monophosphate end resulted in a low but appreciable conversion to a T1 fragment that comigrated with the cyclic phosphate species (Fig. 3, bottom panel, lanes 5 and 6).

The time course of the 2′,3′-cyclic phosphodiesterase (CPDase) activity of RtcB is shown in Fig. 5B. Opening of the cyclic end was relatively rapid and efficient (65% of the input 2′,3′-p substrate was hydrolyzed in 2 min), and the reaction attained an end point with 80% of the cyclic ends opened. The inference that the CPDase reaction is reversible was consistent with the kinetic profile of the reaction of RtcB with a 5′-phosphorylated, 3′-monophosphate strand, wherein we saw a slow but steady conversion of the 3′-phosphate end to a species that comigrated with the 2′,3′-p T1 fragment (Fig. 5D). The extent of this conversion was 15% at 45 min. The RtcB reaction mixtures did not include ATP, which is the requisite high energy substrate for RNA 3′-cyclization catalyzed by bacterial RtcA (24, 25). Given that RtcA gene expression is repressed in E. coli under standard growth conditions (25), and the purity of the recombinant RtcB preparation (Fig. S2), we thought it unlikely that the 3′-phosphate cyclization observed for RtcB could be caused by RtcA contamination. To confirm this idea, we purposefully included 0.1 mM ATP in the RtcB reaction mixtures and examined the kinetic profile of cyclization, assuming that ATP would strongly stimulate the generation of cyclic ends if RtcA was the culprit (25). However, we found no effect of ATP on the rate or yield of cyclic phosphate product (not shown). Thus, we surmise that RtcB has an intrinsic CPDase activity that favors the forward hydrolysis reaction under the conditions of our assays.

The temporal profile of the RtcB intramolecular ligation reaction with the 5′-OH/2′,3′-p 20-mer substrate was consistent with the intermediacy of a 3′-monophosphate end in the sealing pathway (Fig. 5A). To wit, the 3′-phosphate T1 fragment was more abundant than the ligated junction fragment at the 2-min time point, at which the 3′-phosphate and ligated species comprised 27 and 18% of the total radioactivity, respec-
Two-step Pathway of End-joining by RNA Ligase RtcB

Effect of RtcB Mutations on CPDase and Ligase Activities—Previously, we used the crystal structure of *Pyrococcus horikoshii* RtcB (20) to guide a mutational analysis of conserved residues in *E. coli* RtcB that we thought might comprise an active site. A water in the putative architectural RtcB active site, coordinated by the equivalents of essential *E. coli* RtcB side chains Asp-75, Cys-78, and His-168 (19), was regarded as a potential mimic of the enzyme-bound metal cofactor. Here we purified the mutant C78A and H168A RtcB proteins to apparent homogeneity via the same sequence of chromatography steps and gel-filtration used to purify the wild-type RtcB enzyme (supplemental Fig. S2) and then tested the wild type and mutants for their ability to execute the component CPDase and 3′-phosphate ligation steps of the proposed RtcB RNA repair pathway.

An RtcB titration experiment established that the extent of intramolecular sealing of 3′-phosphate and 5′-OH ends was proportional to the concentration of wild-type RtcB from 50 to 500 nM, attaining an end point of 98% end joining at 500–1000 nM (Fig. 7A). By contrast, C78A was inert in 3′-phosphate ligation even at 1000 nM, the highest level of protein tested (Fig. 7A). The H168A mutant displayed 3′-phosphate ligation activity only at 1000 nM input protein (17% of substrate ligated).
Mutational effects on the CPDase reaction are shown in Fig. 7B. Whereas opening of the cyclic phosphate by wild-type RtcB increased in proportion to enzyme concentration and was nearly complete at 500–1000 nM RtcB, the C78A and H168A mutants were virtually inert at up to 1000 nM input protein. We infer that the CPDase and 3'-phosphate ligase reactions are catalyzed by a common active site.

A Role for GTP in 3'-Phosphate/5'-OH Ligation—In our initial characterization of RtcB, we noted that 0.1 mM GTP stimulated the sealing of a broken RNA stem-loop with 2',3'-cyclic phosphate and 5'-OH termini (18). Consequently, we have included GTP in the RtcB reaction mixtures in all experiments presented above. The new findings that RtcB ligates a 3'-phosphate end to a 5'-OH end raised the question of whether GTP might serve as an energy-rich substrate in the ligation reaction, analogous to the way classic RNA ligases and RNA cyclase use ATP for end activation. Thus, we assayed 3'-phosphate ligase activity in the presence and absence of GTP. The extents of product formation are plotted as a function of input RtcB in supplemental Fig. S3, wherein each datum is the average of three separate enzyme titration experiments. We found that RtcB could execute the 3'-phosphate ligation reaction in the absence of GTP; ligation proceeded to near completion at 1000 nM enzyme. However, 3'-phosphate ligase activity was stimulated 3-fold by 0.1 mM GTP (supplemental Fig. S3). By contrast, 0.1 mM GDP had little or no stimulatory effect (supplemental Fig. S3).

If the analogy between RtcB and classic ligases is valid, then one predicts that RtcB would react with GTP to form a covalent enzyme-guanylate intermediate (see below), in which case the observed GTP-independent 3'-phosphate sealing activity would be executed by a population of pre-formed RtcB-GMP (or RtcB-GDP) in the recombinant enzyme preparation. To explore this point further, we modified the purification protocol for recombinant RtcB in an effort to reduce or eliminate any residual nucleotide or metals. To wit: (i) during the second Ni-agarose step, the flow-through fraction containing tag-free RtcB was immediately adjusted to 20 mM EDTA and incubated for 1 h, and (ii) the EDTA-treated RtcB was then purified by gel filtration through a column of Superdex 200 in buffer containing 1 mM EDTA. This new RtcB enzyme preparation was strictly dependent on added manganese and either GTP or dGTP (0.1 mM) for its 3'-phosphate/5'-OH ligation activity (Fig. 8). ATP, CTP, and UTP (0.1 mM) were ineffective in lieu of GTP (Fig. 8).

RtcB Reacts with GTP to Form a Covalent RtcB-guanylate Adduct—Incubation of RtcB with 10 μM [α-32P]GTP in the presence of 2 mM manganese resulted in the transfer of radiolabel to the RtcB polypeptide to form an SDS-resistant adduct that migrated as a single discrete species of ~45 kDa during SDS-PAGE (Fig. 9C). The extent of label transfer was proportional to input RtcB concentration (Fig. 9A); ~10–15% of the added RtcB protomers were 32P-labeled in vitro under these conditions. No label transfer from [γ-32P]GTP to RtcB occurred in the absence of manganese (Fig. 9A).

We found that no label was transferred to RtcB when the enzyme was reacted with 10 μM [γ-32P]GTP (Fig. 9B). Therefore we surmise that the radiolabeled species is most likely a

**FIGURE 7. Effect of RtcB active site mutations on 3'-phosphate ligation and CPDase activities.** A, ligase reaction mixtures (10 μl) containing 50 mM Tris-HCl (pH 8.0), 0.1 mM GTP, 2 mM MnCl2, 20 nM 32P-labeled 20-mer pRNA as shown and 0, 50, 100, 200, 500, or 1000 nM of wild-type (WT) RtcB or mutants C78A or H168A (see supplemental Fig. S2) were incubated at 37 °C for 30 min. The products were digested with RNase T1 and then analyzed by PAGE. B, CPDase reaction mixtures (10 μl) containing 50 mM Tris-HCl (pH 8.0), 0.1 mM GTP, 2 mM MnCl2, 20 mM 32P-labeled 20-mer pRNA as shown and 0, 50, 100, 200, 500, or 1000 nM of WT or mutant RtcB proteins were incubated at 37 °C for 30 min. The products were digested with RNase T1 and then resolved by PAGE.

**Two-step Pathway of End-joining by RNA Ligase RtcB**
the reaction of RtcB with [α-32P]GTP to form the radiolabeled adduct (not shown). Thus, in multiple respects, including metal cofactor specificity, nucleotide specificity, and reliance on Cys-78 in the active site, the properties of the RtcB guanylylation reaction and the RtcB ligation reaction are concordant. Thus, we speculate that the RtcB-GMP complex is a likely intermediate in the 3'-phosphate ligation reaction.

Chemical Sensitivity of the RtcB-guanylate Adduct—The nature of the linkage between RtcB and the 32P-guanylate was evaluated by chemical sensitivity tests (27), whereby the product of the label-transfer reaction was treated with EDTA and SDS and then exposed to acid, alkali, or hydroxylamine. The protein-guanylate adduct was stable to heating for 5 min at 75 °C in 0.15 M NaOH (Fig. 10). By contrast, 89% of the adduct was hydrolyzed by heating for 5 min at 75 °C in 0.15 M HCl, a treatment to which the RtcB polypeptide was itself stable (Fig. 10). The RtcB-[32P]guanylate adduct was eliminated by exposure for 30 min at 37 °C to 0.6 M hydroxylamine (pH 5.0) but was resistant to treatment in parallel with 0.6 M sodium acetate (pH 5.0) (Fig. 10). The chemical lability of the adduct to acid and hydroxylamine and its resistance to alkali are characteristic of a phosphoramidate bond (27). In the case of RtcB, where the active site is surmised from the crystal structure of the Pyrococcus homolog (20) and the mutational analysis inspired thereby (19), we suspect that the guanylate is attached to one of the essential histidine residues.

DISCUSSION

The present study establishes two facts that weight heavily against the one-step direct ligation mechanism depicted in Fig. 1A: (i) RtcB is adept at joining a 3'-monophosphate to a 5'-OH and (ii) RtcB has a CPDase activity that hydrolyzes a 2',3'-cyclic phosphate end to a 3'-monophosphate. The available data favor the two-step pathway of RNA sealing depicted in Fig. 1B, in which the CPDase acts upstream of the 3'-phosphate ligase and both activities are intrinsic to RtcB. The CPDase of RtcB is envisioned as a straightforward in-line attack by water (or hydroxyl ion) on the cyclic phosphate apical to the ribose O2' leaving group. This step is metal-dependent, with manganese being the optimal cofactor. The structure of archaeal RtcB suggests a binding site for one or more metals that is rich in essential histidines and an essential cysteine. The one or more metals could promote CPDase catalysis by coordinating the water/hydroxyl nucleophile, coordinating and promoting expulsion of the ribose 2' oxyanion leaving group, and/or coordinating the non-bridging phosphate oxygens and stabilizing the transition state.

The 3'-phosphate ligation step of the RtcB pathway presents a challenge to phosphoryl transfer orthodoxy. It is unappealing on energetic grounds to suppose that the ligation reaction occurs by the direct attack of the terminal O5' nucleophile on the 3'-monophosphate to yield a 3',5'-phosphodiester, because such a mechanism calls for the expulsion of one of the non-bridging phosphate oxygens (as hydroxide or, eventually, water) while the bridging phosphomonester linkage remains intact. Rather, it is more appealing, and less of a departure from classic ligases, to invoke a mechanism entailing RNA 3' end activation via one or more high energy inter-
Two-step Pathway of End-joining by RNA Ligase RtcB

FIGURE 9. Reaction of RtcB with [α-32P]GTP to form a covalent RtcB-guanylate adduct. A, reaction mixtures (10 μl) containing 50 mM Tris-HCl (pH 8.0), 10 μM [α-32P]GTP, either 2 mM MnCl2 (+Mn) or no added divalent cation (−Mn), and increasing concentrations of RtcB (0.5, 1, 2.5, and 5 μM, proceeding from left to right in each titration series) were incubated at 37 °C for 10 min. B, reaction mixtures (10 μl) containing 50 mM Tris-HCl (pH 8.0), either 10 μM [α-32P]GTP, 10 μM [γ-32P]GTP, or 10 μM [α-32P]ATP as specified (at equivalent specific radioactivity), and either 5 μM RtcB plus 2 mM MnCl2 or 5 μM chloroŭea virus DNA ligase (ChV-Lig) plus 5 mM MgCl2 were incubated at 37 °C for 10 min. C, reaction mixtures (10 μl) containing 50 mM Tris-HCl (pH 8.0), 10 μM [α-32P]GTP, 5 μM RtcB, and 2 mM of the indicated divalent cation (as the chloride salt) were incubated at 37 °C for 10 min. D, reaction mixtures (10 μl) containing 50 mM Tris-HCl (pH 8.0), 10 μM [α-32P]GTP, 5 μM RtcB, 2 mM MnCl2, and 2 μM of the indicated divalent cation (as the chloride salt) were incubated at 37 °C for 10 min. The reactions were quenched by adding 10 μl of a solution containing 100 mM Tris (pH 6.8), 10 mM EDTA, 200 mM DTT, 4% SDS, 20% glycerol, 0.2% bromphenol blue. The mixtures were then analyzed by SDS-PAGE. Autoradiographs of the dried gels are shown. The positions and sizes of pre-stained marker proteins are indicated on the left.

mediates. Indeed, we found here that 3′-phosphate ligation by RtcB depends on GTP and that RtcB reacts with GTP in the absence of nucleic acid to form a covalent RtcB-guanylate adduct. These properties are reminiscent of the many classic polynucleotide ligases that act via covalent ligase-adenylate intermediates. Extending the analogy further, we speculate that RtcB-guanylate is the first of two activated ligation reaction intermediates, the second being RNA(3′)pp(5′)G generated by transfer of guanylate from RtcB to the RNA 3′-monophosphate terminus. Guanylation of the RNA 3′-phosphate would activate it for attack by an RNA 5′-OH end to form the sealed phosphodiester and liberate GMP. This mechanism echoes how RNA cyclase enzymes transfer AMP from a covalent enzyme-adenylate intermediate to an RNA 3′-phosphate to form RNA(3′)pp(5′)A, thereby activating the 3′-phosphate for an attack by the vicinal ribose O2′ to yield a 2′,3′-cyclic phosphate and liberate AMP (24, 28). In this vein, we speculate that the apparent reversal of the CPDase reaction of RtcB (seen when RNA ligation is prevented by a 5′-phosphate) reflects the use of the ribose O2′ as a nucleophile (instead of an RNA 5′-OH end) for attack on RNA(3′)pp(5′)G.

Additional studies will be required to map the site of covalent RtcB guanylation, validate the intermediacy of RtcB-GMP, and capture the putative (and presumably evanescent) guanylated RNA intermediate. A complete mechanistic picture will ultimately hinge on obtaining cocrystal structures of RtcB in complexes with manganese, GTP, GMP, and a 3′-phosphate RNA substrate.

Finally, the two-step RtcB pathway in Fig. 1B has important implications for the spectrum of RNA repair reactions that RtcB might perform. Specifically, our findings indicate that the substrates for sealing by RtcB need not be limited to RNAs with 2′,3′-p ends. RNA-damaging ribotoxins and tRNA-splicing endonucleases characteristically generate 2′,3′-p termini as their end-products, and these broken RNAs have been considered the most plausible substrates for splicing or repair by RtcB ligases. However, many other ribonucleases generate 3′-monophosphates, which can now be considered potential RtcB substrates. Indeed, our finding of an intrinsic 3′-phosphate ligation activity in E. coli RtcB provides a plausible explanation for the results of Popow et al. (17), who partially purified a ligase complex containing mammalian RtcB from a HeLa cell extract based on its ability to join a 3′-monophosphate end to a 5′-OH end. These investigators documented by mass spectrometry the presence of human RtcB in the ligase preparation, but they did not detect the presence of the human RNA 3′-phosphate cyclase protein (Rtc1). Based on our findings, we speculate that human RtcB may also exploit a two-step reaction pathway.
REFERENCES

1. Konarska, M., Filipowicz, W., and Gross, H. J. (1982) Proc. Natl. Acad. Sci. U.S.A. 79, 1474–1478
2. Greer, C. L., Peebles, C. L., Gegenheimer, P., and Abelson, J. (1983) Cell 32, 537–546
3. Amitsur, M., Levitz, R., and Kaufmann, G. (1987) EMBO J. 6, 2499–2503
4. Sidrauski, C., Cox, J. S., and Walter, P. (1996) Cell 87, 405–413
5. Gonzalez, T. N., Sidrauski, C., Dörfler, S., and Walter, P. (1999) EMBO J. 18, 3119–3132
6. Nandakumar, J., Schwer, B., Schaffrath, R., and Shuman, S. (2008) Mol. Cell 31, 278–286
7. Schwer, B., Sawaya, R., Ho, C. K., and Shuman, S. (2004) Proc. Natl. Acad. Sci. U.S.A. 101, 2788–2793
8. Nandakumar, J., Shuman, S., and Lima, C. D. (2006) Cell 127, 71–84
9. Sawaya, R., Schwer, B., and Shuman, S. (2003) J. Biol. Chem. 278, 43928–43938
10. Wang, L. K., Schwer, B., Englert, M., Beier, H., and Shuman, S. (2006) Nucleic Acids Res. 34, 517–527
11. Martins, A., and Shuman, S. (2005) RNA 11, 1271–1280
12. Chan, C. M., Zhou, C., and Huang, R. H. (2009) Science 326, 247
13. Filipowicz, W., and Shatkin, A. J. (1983) Cell 32, 547–557
14. Laski, F. A., Fire, A. Z., RajBhandary, U. L., and Sharp, P. A. (1983) J. Biol. Chem. 258, 11974–11980
15. Zofallova, L., Guo, Y., and Gupta, R. (2000) RNA 6, 1019–1030
16. Englert, M., Sheppard, K., Aslanian, A., Yates, J. R., 3rd, and Söll, D. (2011) Proc. Natl. Acad. Sci. U.S.A. 108, 1290–1295
17. Popow, J., Englert, M., Weitser, S., Schleiffer, A., Mierzwia, B., Mechtler, K., Trowitzsch, S., Will, C. L., Lührmann, R., Söll, D., and Martinez, J. (2011) Science 331, 760–764
18. Tanaka, N., and Shuman, S. (2011) J. Biol. Chem. 286, 7727–7731
19. Tanaka, N., Meineke, B., and Shuman, S. (2011) J. Biol. Chem. 286, 30253–30257
20. Okada, C., Maegawa, Y., Yao, M., and Tanaka, I. (2006) Proteins 63, 1119–1122
21. Shuman, S. (1998) Mol. Cell 1, 741–748
22. Wang, L. K., and Shuman, S. (2002) Nucleic Acids Res. 30, 1073–1080
23. Wang, L. K., Ho, C. K., Pei, Y., and Shuman, S. (2003) J. Biol. Chem. 278, 29454–29462
24. Tanaka, N., Smith, P., and Shuman, S. (2010) Structure 18, 449–457
25. Genschik, P., Drabikowski, K., and Filipowicz, W. (1998) J. Biol. Chem. 273, 25516–25526
26. Nair, P. A., Nandakumar, J., Smith, P., Odell, M., Lima, C. D., and Shuman, S. (2007) Nat. Struct. Mol. Biol. 14, 770–778
27. Gumport, R. I., and Lehman, I. R. (1971) Proc. Natl. Acad. Sci. U.S.A. 68, 2559–2563
28. Filipowicz, W., Strugala, K., Konarska, M., and Shatkin, A. J. (1985) Proc. Natl. Acad. Sci. U.S.A. 82, 1316–1320

Two-step Pathway of End-joining by RNA Ligase RtcB

FIGURE 10. Chemical stability of the RtcB-[32P]guanylate adduct. Reaction mixtures (140 μl) containing 50 mM Tris-HCl (pH 8.0), 10 mM [α-32P]GTP, 2 mM MnCl2, and 5 μM RtcB were incubated at 37 °C for 1 min. The mixtures were then adjusted to 0.5% SDS and 5 mM EDTA. Aliquots (20 μl) were withdrawn and either received no further additives (lane –) or were adjusted to 600 mM hydroxylamine (pH 5.0) or 600 mM sodium acetate (pH 5.0), and then incubated at 37 °C for 30 min. Additional aliquots (20 μl) were withdrawn and adjusted to either 150 mM Tris-HCl (pH 7.4), 150 mM HCl, 150 mM NaOH, or 150 mM NaCl; these samples were heated at 75 °C for 5 min; the HCl-treated sample was neutralized by addition of NaOH to 150 mM and the NaOH-treated sample was neutralized by adding HCl to 150 mM. The samples were supplemented with 20 μl of a solution containing 100 mM Tris (pH 6.8), 10 mM EDTA, 200 mM DTT, 4% SDS, 20% glycerol, 0.2% bromophenol blue. One-half of each sample was analyzed by SDS-PAGE, followed by autoradiography of the dried gel to visualize the RtcB-[32P]guanylate adduct (top panel). The other half of each sample was analyzed by SDS-PAGE, and the gel was stained with Coo massie Blue dye to visualize the RtcB polypeptide (bottom panel); the positions of 50- and 37-kDa pre-stained marker proteins are indicated on the left.