Transcript assisted phosphodiester bond hydrolysis by eukaryotic RNA polymerase II

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Hydrolysis of the phosphodiester bonds of the transcript by bacterial RNA polymerase is assisted by 3′NMP of the RNA. Here we provide evidence that this mechanism is also involved in RNA cleavage by eukaryotic RNA polymerase II, suggesting that transcript assisted hydrolysis has emerged before divergence of bacteria and archaea/eukaryotes.

Multisubunit RNA polymerase (RNAP), the enzyme accomplishing transcription in all living organisms, has emerged before the divergence of bacteria and archaea/eukaryotes. Accordingly, the molecular mechanisms involved in RNA synthesis are highly conserved in evolution. Besides the synthesis of phosphodiester bonds (and pyrophosphorolysis, which is a direct reversal of this reaction) during transcript elongation, RNAP active center can catalyze the hydrolysis of the phosphodiester bonds of the nascent RNA. As well as the synthesis, the hydrolysis is catalyzed by the two metal (Mg²⁺) ion mechanism. In addition, in bacterial RNAP, a flexible domain of the active center, the Trigger Loop, plays a critical role in hydrolysis by participating in the reaction as a general base.

This appears to be different from eukaryotic RNA polymerase II (RNAP II), whose Trigger Loop fails to adopt a catalytically active conformation during hydrolysis, explaining a much slower intrinsic hydrolysis by RNAP II as compared with bacterial RNAPs.

Earlier, we have discovered that an NMP at the 3′ end of the RNA transcript in the bacterial elongation complex also participates in catalysis of phosphodiester bond hydrolysis. For this to occur, RNAP has to backtrack by 1 base pair, thus positioning the penultimate (second from the 3′ end of the RNA) phosphodiester bond in the active center, making it ready for cleavage. The 3′NMP disengages from the template strand and flips backward to approach the site of the reaction, and helps to chelate the second Mg²⁺ ion, Mg²⁺II (which otherwise is bound weakly), to position the attacking water molecule and, possibly, to participate in the catalysis as a general acid/base. Such transcript-assisted hydrolysis of the second phosphodiester bond becomes even more prominent when RNAP is stabilized in the 1 base pair backtracked state via misincorporation of NMP that is not complementary to the base in the template strand. Chemical groups of 3′AMP, CMP, GMP and UMP contribute differently during the hydrolysis. Furthermore, the nature of the misincorporation event, i.e., the base in the template strand, may also influence the involvement of the chemical groups of the 3′NMP in the reaction. The erroneously incorporated NMP can be imagined to help to excise itself from the transcript, thus contributing to the proofreading of misincorporated events, i.e., overall fidelity of transcription.

The “self-correcting” function of the transcript led to the proposition that transcript assisted second phosphodiester bond hydrolysis could be an ancient feature of transcription by multisubunit RNAPs, which may have emerged before the divergence of bacterial and archaeal/eukaryotic lineages. A possible involvement of the 3′NMP in the RNA hydrolysis by eukaryotic RNAP became evident from the crystal structure of the backtracked elongation complex of RNAP II.

Keywords: RNA polymerase II, transcript assisted proofreading, phosphodiester bond hydrolysis, elongation complex, backtracking

Abbreviations: RNAP, RNA polymerase; RNAP II, RNA polymerase II; NMP, nucleotide monophosphate; NTA, nitrilotriacetate

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As was proposed for bacterial RNA polymerase II, the 3′NMP (GMP in this case) flips backward coming closer to the second phosphodiester bond and Mg²⁺, possibly making itself available for assisting the reaction. This was also consistent with the faster cleavage by RNA polymerase II of the second phosphodiester bond compared with the ultimate one, even in the complexes that were not stabilized in the backtracked state. However, no biochemical evidence for the involvement of the 3′NMP, i.e., transcript assistance, in the hydrolysis of the second phosphodiester bond exists so far.

We decided to analyze the possibility of the assistance from the transcript’s 3′NMP during second phosphodiester bond hydrolysis by RNA polymerase II active center. To do so, we used artificial elongation complexes assembled with *S. cerevisiae* RNA polymerase II, fully complementary synthetic template and non-template DNA strands and synthetic RNA transcript (see scheme in Fig. 1B). These complexes are indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences. The complexes were indistinguishable from “native” stalled elongation complexes obtained by transcription on the double-stranded DNA from the promoter, but allow omitting the complicated step of transcription initiation and easy changing of the transcribed sequences.
We misincorporated GMP, UMP and AMP opposite to dTMP, dTMP and dGMP in the template strand, respectively, and analyzed RNA hydrolysis in the formed misincorporated elongation complexes (mEC15G, mEC15U and mEC15A) (Fig. 1B). We determined the affinity to Mg\(^{2+}\)II (Km[Mg\(^{2+}\)]) and the rate of the reaction in the saturating Mg\(^{2+}\) (k\(_{\text{cat}}\)). As seen from Figure 1C, the maximal rate of the second phosphodiester bond hydrolysis in mEC15A was ~4 times slower than that in mEC15G or mEC15A, suggesting that 3’purine increases the rate of the reaction as compared with pyrimidines. Km[Mg\(^{2+}\)] in mEC15G was 3~4 times higher than that in mEC15G or mEC15A. This result suggests that the 3’ UMP and GMP may participate in the chelation of Mg\(^{2+}\)II, while 3’ AMP may not provide additional coordination bonds.

To analyze if chemical groups of 3’purine are involved in the accelerated second phosphodiester bond hydrolysis, we used chemically modified 3’AMP. We misincorporated α-thio-AMP, 7-deaza-AMP and N6-methyl-AMP at the 3’ end of the transcript and measured the rate of second phosphodiester bond hydrolysis in the resultant complexes. Given that 3’AMP (i.e., its chemical groups) unlikely participates in the Mg\(^{2+}\) binding, we measured the rate of the reaction only in 10 mM Mg\(^{2+}\). As seen from Figure 1D, N6-methyl group had little effect on the rate of the reaction. However, substitution of nitrogen in the position 7 of the purine rings with carbon reduced the rate of the reaction ~3-fold, while the thio-group in the ultimate phosphodiester bond slowed down the reaction ~7-fold.

The above result suggests that the nitrogen in position 7 of the adenine ring may participate in the penultimate phosphodiester bond hydrolysis as an acid/base and/or by coordinating the attacking water molecule. This hypothesis is supported by the crystal structure of the backtracked RNAP II elongation complex (pdb: 3GTJ), in which the N7 of the 3’guanine moiety is turned toward the penultimate phosphodiester bond where it can assist the reaction (Fig. 1A). The phosphate of the 3’GMP in this structure is also positioned so that it can directly or through the network of hydrogen bonding interact with the attacking water molecule (Fig. 1A), which is consistent with our results.

All together, our results suggest that, like bacterial RNAP, RNAP II also may use the transcript-assisted hydrolysis of the phosphodiester bond. As mentioned above, the Trigger Loop of RNAP II seems not to fold fully in the backtracked elongation complex, and thus cannot participate in the transcript assisted hydrolysis, as it does in the case of bacterial RNAP.\(^2\) This deviation and some differences in the amino acid content of the active centers of bacterial RNAP and RNAP II likely influence the way the chemical groups of 3’NMPs are involved in the reaction. For example, N7 of the 3’AMP of bacterial elongation complex was proposed to participate in the chelation of Mg\(^{2+}\)II.\(^3\) In the case of RNAP II, however, 3’AMP does not contribute strongly to binding of Mg\(^{2+}\)II. Instead, N7 of 3’AMP seems to accelerate the reaction rate. In contrast, however, for example, the phosphate group of 3’AMP seems to be involved in the reaction similarly for both, bacterial RNAP\(^4\) and RNAP II.

The 3’AMP assisted hydrolysis is ~4 times faster than 3’UMP assisted hydrolysis in both bacterial and RNAP II elongation complexes. It is also ~4 times faster than cleavage assisted by 3’CMP in bacterial RNAP (we did not analyze assistance from 3’CMP for RNAP II). Though cleavage assisted by 3’GMP in bacterial elongation complex is ~4~5 times slower than with 3’AMP,\(^5\) the rates of cleavage by RNAP II assisted by 3’AMP or 3’GMP are close. Together, these data may indicate that the 3’purine accelerates phosphodiester bond hydrolysis more efficiently than 3’pyrimidines, possibly due to the presence of the imidazole ring. It is possible, that the participation of 3’purines in the reaction has been similar earlier in evolution but diverged later, with N7 of 3’AMP losing its ability to chelate Mg\(^{2+}\) during cleavage by RNAP II, and N7 of 3’GMP losing the ability to accelerate the reaction by bacterial RNAP.

Deeper biochemical analysis, such as determining pH profiles of reactions and analysis of a more diverse panel of chemically modified 3’NMPs are required to build the full picture of similarities/differences of transcript assisted RNA cleavage between bacterial and eukaryotic RNAPs. However, the above data together with our earlier results\(^6\) support an intriguing hypothesis that transcript-assisted phosphodiester bond hydrolysis may have emerged before the divergence of bacteria and archaea/eukaryotes, at the stage of the Last Universal Common Ancestor (LUCA). This mechanism may have been critical in the absence of elongation cleavage factors (evolutionary unrelated factor S of archaea/eukaryotes and Gre factor of bacteria, which must have emerged after divergence of domains of life) to rescue backtracked and/or misincorporated transcription elongation complexes.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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