Wright, Jaden R.

2011

Pre-steady-state kinetic analysis of the three Escherichia coli pseudouridin synthases TruB, TruA, and RluA reveals uniformly slow catalysis

Department of Chemistry and Biochemistry

https://hdl.handle.net/10133/5384

Downloaded from OPUS, University of Lethbridge Research Repository
Pre-steady-state kinetic analysis of the three *Escherichia coli* pseudouridine synthases TruB, TruA, and RluA reveals uniformly slow catalysis

JADEN R. WRIGHT, LAURA C. KEFFER-WILKES, SELINA R. DOBING, and UTE KOTHE

Department of Chemistry and Biochemistry, University of Lethbridge, Lethbridge, Alberta, Canada T1K 3M4

ABSTRACT

Pseudouridine synthases catalyze formation of the most abundant modification of functional RNAs by site-specifically isomerizing uridines to pseudouridines. While the structure and substrate specificity of these enzymes have been studied in detail, the kinetic and the catalytic mechanism of pseudouridine synthases remain unknown. Here, the first pre-steady-state kinetic analysis of three *Escherichia coli* pseudouridine synthases is presented. A novel stopped-flow absorbance assay revealed that substrate tRNA binding by TruB takes place in two steps with an overall rate of 6 sec\(^{-1}\). In order to observe catalysis of pseudouridine formation directly, the traditional tritium release assay was adapted for the quench-flow technique, allowing, for the first time, observation of a single round of pseudouridine formation. Thereby, the single-round rate constant of pseudouridylation (\(k_C\)) by TruB was determined to be 0.5 sec\(^{-1}\). This rate constant is similar to the \(k_{cat}\) obtained under multiple-turnover conditions in steady-state experiments, indicating that catalysis is the rate-limiting step for TruB. In order to investigate if pseudouridine synthases are characterized by slow catalysis in general, the rapid kinetic quench-flow analysis was also performed with two other *E. coli* enzymes, RluA and TruA, which displayed rate constants of pseudouridine formation of 0.7 and 0.35 sec\(^{-1}\), respectively. Hence, uniformly slow catalysis might be a general feature of pseudouridine synthases that share a conserved catalytic domain and supposedly use the same catalytic mechanism.

Keywords: pseudouridine; pseudouridine synthase; kinetic mechanism; RNA modification; pre-steady-state kinetic analysis

INTRODUCTION

Functional RNAs are typically modified within the cell to enhance their structural and functional properties. Numerous different modifications of the ribose and the nucleobases have been identified (Limbach et al. 1994), of which pseudouridines (Ψ) are the most abundant modifications found in all major functional RNAs such as tRNA, rRNA, snRNA, and snoRNA (Charette and Gray 2000). Pseudouridines are isomers of uridines differing only in the glycosidic bond, which is a C5-glycosidic bond in pseudouridine instead of the canonical N1-glycosidic bond (Fig. 1A). While pseudouridines can form the same base-pairing interactions as uridines, the extra hydrogen bond donor in the nitrogenous base, the N1H imino group, can form additional interactions. For example, it has been shown that the N1H imino group can stabilize a water molecule between the nucleobase and the phosphate backbone, thereby rigidifying the local RNA structure (Arnez and Steitz 1994). Besides their widely accepted structural role, pseudouridines may also play additional functional roles for rRNA and snRNA. Most pseudouridines are located close to the functional centers of the RNAs, such as the ribosomal peptidyltransferase center, the ribosomal decoding center, or the spliceosomal branch site (Charette and Gray 2000; Decatur and Fournier 2002). Specifically, pseudouridines have been proposed to play a role in translation termination (Ejby et al. 2007; Liang et al. 2007) and in the first step of splicing by positioning the branch site adenosine (Newby and Greenbaum 2002; Yang et al. 2005). In accordance with their importance for RNA structure and function, the abundance of pseudouridines increases with the evolutionary complexity of an organism; while *Escherichia coli* rRNA contains 11 pseudouridines, about 100 of these modifications have been identified in human rRNA (Ofengand 2002).
The formation of pseudouridines is catalyzed by enzymes called pseudouridine synthases, which can be grouped in five families found in bacteria plus one additional pseudouridine synthase found in archaea and some eukaryotes (Hamma and Ferré-D’Amáré 2006; McCleverty et al. 2007). Each bacterial pseudouridine synthase catalyzes pseudouridine formation at one or a few distinct sites within cellular RNA. In addition, eukaryotes and archaea use H/ACA small (nucleolar) ribonucleoproteins which catalyze pseudouridylation at many different sites within cellular RNA with the help of many different box H/ACA guide RNAs (Ye 2007). Within the last decade, crystal structures of pseudouridine synthases from all six families have been determined (Gu et al. 1998; Wrzesinski et al. 1995; Raychaudhuri et al. 1999; Hoang et al. 2006). While the basic steady-state kinetic parameters of many pseudouridine synthases have been determined (Gu et al. 1998; Huang et al. 1998; Ramamurthy et al. 1999b), not much insight has been gained into the mechanism of pseudouridine synthases, which has been suggested to consist of up to six steps (Arluison et al. 1999). So far, only some information on the kinetics of substrate tRNA binding by the Saccharomyces cerevisiae pseudouridine synthase Pus1 has been reported (Arluison et al. 1999).

Here, we have analyzed for the first time the kinetic mechanism of pseudouridine formation using the model enzyme TruB with the aim of identifying the rate-limiting step during pseudouridylation. For this purpose, two novel rapid kinetic assays were established, enabling us to monitor
RESULTS AND DISCUSSION

Absorbance assay for tRNA binding to TruB

In order to dissect the kinetic mechanism of pseudouridylation, the well-studied E. coli pseudouridine synthase TruB was used as a model enzyme (Nurse et al. 1995). TruB was overexpressed containing an N-terminal histidine-tag and purified by affinity and size-exclusion chromatography, similarly to previous reports (Nurse et al. 1995; Hoang and Ferré-D’Amare 2001). For binding measurements, the active site residue aspartate 48 in TruB was mutated to asparagine (Ramamurthy et al. 1999a; Hoang et al. 2005). This substitution completely abolishes any catalytic activity as confirmed by a tritium release assay detecting pseudouridine formation (data not shown) and thus prevents tRNA modification and product release while retaining the ability to bind tRNA. First, the interaction of E. coli tRNA^Phe with TruB D48N was characterized by nitrocellulose filtration. For this purpose, [3H]-labeled tRNA^Phe was generated by in vitro transcription, purified by anion exchange chromatography, and incubated with increasing amounts of TruB D48N. tRNA^Phe bound to the enzyme remained on the nitrocellulose membrane during filtration and was quantified by scintillation counting (Fig. 2A). Fitting of the binding curve to a hyperbolic equation revealed a dissociation constant, K_D, for the interaction of TruB D48N with tRNA^Phe of 1.4 ± 0.3 µM. This value is in excellent agreement with previously published results for TruB D48A and TruB D48C determined by nitrocellulose filtration (Ramamurthy et al. 1999a), thus confirming the quality of our protein and tRNA preparations.

In order to study the kinetic mechanism of the TruB–tRNA interaction, we developed a new absorbance-based assay for subsequent use in rapid-kinetic stopped-flow measurements. As in the nitrocellulose filtration assay, unlabeled, in vitro transcribed and purified tRNA^Phe was incubated with increasing concentrations of TruB D48N. This time, the absorbance of the reaction mixture was recorded at 260 nm. In parallel, TruB D48N was titrated into buffer; the increasing absorbance of this solution was also monitored and subtracted from the absorbance of the reaction mixture. Thereby, only the absorbance change due to tRNA binding to TruB D48N was observed. The absorbance of tRNA^Phe increases upon binding to TruB D48N (Fig. 2B); this hyperchromic effect might be due to a conformational change in the tRNA upon binding TruB, resulting in a larger proportion of unstacked bases in the tRNA. As in the nitrocellulose filtration assay, a hyperbolic binding curve is observed, and fitting yielded a dissociation constant of 0.34 ± 0.06 µM. Interestingly, this value is significantly lower than the K_D determined by nitrocellulose filtration, which can be explained by the different nature of these assays. In contrast to the nitrocellulose filtration assay, the absorbance assay is an equilibrium method where both bound and unbound tRNA^Phe are present in the analyzed sample. This assay should, therefore, provide the true dissociation constant. In the nonequilibrium filtration assay, however, the washing step separates bound from unbound tRNA^Phe. While washing was performed fast and with precooled buffer, it is likely that some TruB D48N–tRNA complex dissociates during this step, thus resulting in a slightly higher K_D. This is also consistent with our observation, as well as with previous reports, that only ~60% binding could be observed in the filter binding assay (Ramamurthy et al. 1999a).

Rapid kinetic stopped-flow analysis of tRNA binding to TruB

To analyze the kinetics of tRNA binding to TruB, 0.75 µM tRNA was rapidly mixed with at least a threefold excess of TruB in a stopped-flow apparatus, and the absorbance at 260 nm was monitored on a msec-to-second time scale. Based on the determined dissociation constant, relatively high tRNA and TruB concentrations were used in order to ensure binding of each tRNA by TruB such that a maximal signal is obtained. Furthermore, due to the excess of enzyme under these conditions, TruB wild-type (WT) can undergo only a single round of catalysis which simplifies the kinetic analysis. As expected from the equilibrium measurements, rapid mixing of tRNA^Phe with TruB D48N resulted in an absorbance increase which took place within 0.5 sec (Fig. 2C). The observed absorbance change could be fitted to a one-exponential equation resulting in an apparent rate, k_app, of 12.7 sec⁻¹. When tRNA^Phe is rapidly mixed with buffer instead of TruB, no absorbance change is observed (Fig. 2C). Next, we performed the same experiment with TruB WT, and again an increase in absorbance was observed within 0.5 sec. Interestingly, the absorbance remained high for >60 sec (data not shown), suggesting that the tRNA may not dissociate from TruB wt after formation of pseudouridine under these conditions. This is in agreement with the comparatively high concentrations used in this assay and with previous findings indicating that TruB WT has a rather high affinity for its product (Ramamurthy et al. 1999a). The time course for tRNA
binding to TruB WT closely resembled the time courses observed with TruB D48N, which shows that the absorbance change is a result of an early event in the interaction of TruB and tRNA which takes place before catalysis.

Fitting to a one-exponential equation revealed an apparent rate, $k_{app}$, of 5.2 sec$^{-1}$ for TruB WT, which is in a similar order of magnitude as the rate observed with TruB D48N. Interestingly, no change in absorbance was observed upon binding of tRNAPhe to the pseudouridine synthase RluA, which targets the anticodon loop (data not shown).

In order to further characterize the kinetics of TruB's interaction with tRNAPhe, similar stopped-flow experiments were carried out using increasing concentrations of TruB WT (2.5–10.0 μM final concentrations). Very similar time courses were obtained as before (data not shown). The amplitudes of the absorbance increases remained constant (data not shown), which is expected as complete binding of all tRNAs occurs at these TruB concentrations, given a $K_D$ for the interaction of ~0.34 μM (vide supra). As before, the time courses were fitted to a one-exponential equation to obtain the apparent rates, $k_{app}$. Interestingly, the apparent rates stayed constant over the TruB concentration range tested, with an average rate of 6.0 ± 1.8 sec$^{-1}$ (Fig. 2D).

This lack of a concentration dependence is in contrast to the linear concentration dependence expected for a bimolecular binding reaction: a faster rate of binding is expected at higher concentrations. Therefore, the absorbance assay apparently monitors a different step than the initial contact of tRNA and TruB. This finding suggests that the interaction of TruB and tRNA occurs in a two-step binding mechanism (Fersht 1998).

These observations are compatible with two different two-step kinetic mechanisms describing the interaction of TruB with its substrate RNA. In a two-step equilibrium reaction, the apparent rate can either increase hyperbolically or decrease hyperbolically with the enzyme concentration, depending on whether the first or the second step is fast, respectively (Fersht 1998). In other words, the absorbance change could result from a slow and rate-limiting conformational change in the unbound tRNA which has to precede rapid binding to TruB. Alternatively (and maybe more likely), the initial and fast encounter of TruB and tRNA could...
occur without a change in absorbance but be followed by a slower conformational change in the tRNA, which is reflected in the increased absorbance. Since no concentration dependence of the apparent rate of binding is observed for TruB, both models are in accordance with the data presented here. No concentration dependence was observed due to the experimental conditions which result in saturated apparent rates at the TruB concentrations used. To distinguish between these models, measurements at lower concentrations of TruB would be necessary; however, this is difficult since the enzyme has to be in excess over tRNA to maintain pseudo-first-order conditions. Accordingly, the tRNA concentration would also have to be reduced, which is not feasible since the absorbance change is small and cannot be detected at lower tRNA concentrations. In summary, we conclude from the presented experiments that substrate binding to TruB takes places with an overall rate of 6 sec$^{-1}$ without assigning this rate to a specific step.

It is tempting to speculate that the observed absorbance increase is a result of conformational changes in the tRNA resulting in unstacking of bases. By comparing the crystal structures of tRNA alone with the reported TruB-RNA complex (Hoang and Ferré-D’Amare´ 2001; Pan et al. 2003), it becomes evident that several conformational changes of the tRNA are required in order to allow it to bind productively to TruB. First, the interaction of the D and TCC arm in the elbow region of tRNA has to be disrupted in order to bind the TCC arm to TruB. Furthermore, three bases flip out of the TCC loop into the catalytic pocket of TruB. Either of these two conformational changes or other tRNA rearrangements could contribute to the absorbance increase observed upon the interaction of tRNA with TruB. However, it is more likely that the partial unfolding of the D and TCC arm contributes mostly to the absorbance change, since RluA also flips out bases into its catalytic pocket (Hoang et al. 2006), but no absorbance change could be recorded for this interaction.

**Kinetic analysis of pseudouridine formation by TruB**

In order to analyze the kinetics of TruB-catalyzed pseudouridylation, we conducted steady-state and pre-steady-state kinetic experiments. For these studies, full-length *E. coli* tRNA$^\text{Phe}$ transcripts were used that contained tritium labels at position C5 of all uracils allowing for the detection of pseudouridylation by the release of tritium upon formation of the C5-glycosidic bond in pseudouridine (Fig. 1A; Cortese et al. 1974). First, we confirmed that our purified components are fully active by assessing the steady-state kinetics of pseudouridine formation by measuring the initial rates of pseudouridine formation by TruB at different substrate tRNA concentrations (Fig. 3A). Our experimental conditions are similar to previous studies, and TruB displayed a $k_{\text{cat}}$ value of 0.7 ± 0.1 sec$^{-1}$ and a $K_{M}$ value of 550 ± 150 × 10$^{-9}$ M, similar to published results (Gu et al. 1998; Wright et al. 2078 RNA, Vol. 17, No. 12

**FIGURE 3.** Michaelis-Menten and quench-flow analysis of pseudouridine formation by TruB. (A) Michaelis-Menten experiment of TruB. Ten nM of enzyme was mixed with increasing concentrations of [3H]-labeled tRNA$^\text{Phe}$, and pseudouridine formation was detected using the tritium release assay. The dependence of the initial rates $v_0$ on the tRNA concentration was fitted to the Michaelis-Menten equation yielding a Michaelis constant, $K_{M}$, of 550 ± 150 nM and a catalytic constant, $k_{\text{cat}}$, of 0.7 ± 0.2 sec$^{-1}$ for TruB. (B) Time courses of pseudouridine formation by TruB under single-round, pre-steady-state conditions. [3H]-labeled tRNA$^\text{Phe}$ (1 μM final concentration) was rapidly mixed with TruB (circles: 2.5 μM, squares: 10 μM final concentration) in a quench-flow apparatus. The percentage of pseudouridine formed at a certain time point was determined using the modified tritium release assay. The apparent rate of pseudouridine formation for each TruB concentration was determined by fitting the time courses to a one-exponential function (smooth lines). (C) Dependence of the apparent rate of pseudouridine formation under single-round conditions on the enzyme concentration. The average apparent rate is 0.5 ± 0.2 sec$^{-1}$ for TruB, as indicated by the horizontal line.
Uniform slow catalysis by pseudouridine synthases

Ramamurthy et al. 1999b). Next, the tritium release assay was adapted to allow for detection of the pre-steady-state kinetics of pseudouridine formation by the quench-flow technique. Upon rapid mixing of tritium-labeled tRNA with an excess of TruB, i.e., under single-turnover conditions, the time courses of pseudouridine formation were followed at increasing TruB concentrations (Fig. 3B). Under these conditions, the tritium release assay monitors the appearance of enzyme-product complex as the tritium is released during the last catalytic step, the formation of the new C-C glycosidic bond. The TruB active site is accessible to water (Hoang and Ferré-D’Amare’ 2001), and therefore, it is conceivable that the released tritium can easily escape the enzyme-product complex before product release occurs. Furthermore, the enzyme is denatured by quenching the reaction with 0.1 M HCl, which further facilitates release of tritium into the supernatant. Therefore, this assay detects the product as soon as it appears in the enzyme-product complex. In the quench-flow experiments, close to 100% pseudouridine formation was detected within 5 sec for all TruB concentrations tested (2.5–10 μM). Single-exponential fitting yielded the apparent rate of pseudouridylation by TruB (Fig. 3C). In the analyzed concentration range, the apparent rate of pseudouridylation was independent of the TruB concentration with an average rate of 0.5 ± 0.2 sec⁻¹. The absence of a concentration dependence suggests that pseudouridine formation by TruB is not limited by tRNA binding under these conditions. Moreover, the rate of substrate binding at 20°C is 6 sec⁻¹ and will be even higher at 37°C, where the tritium release assays were conducted, thus further supporting the finding that tRNA binding is not rate-limiting for TruB. In conclusion, the rate of pseudouridine formation measured here directly reflects the rate constant of pseudouridine catalysis (kₚ = 0.5 ± 0.2 sec⁻¹). Here, pseudouridine catalysis is understood as the overall process comprising glycosidic bond cleavage, base rotation, and new C-C glycosidic bond formation, as these steps cannot be distinguished by the tritium release assay. It cannot be excluded that kₚ also comprises flipping of the target U55 into the active site of TruB if this flipping would be substantially slower than the conformational changes observed in the absorbance experiments.

Notably, this rate constant of pseudouridine formation, kₚ, determined under single-round conditions is very similar, within the experimental error, to the k_cat obtained from steady-state experiments under multiple-turnover conditions. The main difference between these experimental conditions is the fact that, in the steady-state experiments, each enzyme has to release the product RNA prior to catalyzing pseudouridine formation in a new tRNA substrate. Therefore, the steady-state experiments in conjunction with the pre-steady-state quench-flow experiments show that the rate-limiting step within the kinetic mechanism of pseudouridine formation by TruB is the catalytic step, and not product release. This is the first time that a detailed kinetic study of a pseudouridine synthase provides important insight into several steps of the kinetic mechanism, i.e., into substrate binding, catalysis, and product release.

Kinetic analysis of pseudouridylation by RluA and TruA

In order to address the question whether catalysis of pseudouridylation is generally a slow step for pseudouridine synthases, we also performed a pre-steady-state kinetic analysis of the E. coli pseudouridine synthases RluA and TruA, representing two pseudouridine synthase families different from TruB. Both enzymes catalyze pseudouridylation in the anticodon arm of many tRNAs, including tRNA_Phe. Specifically, RluA modifies position 32, and TruA targets position 39, in tRNA_Phe (Fig. 1B; Turnbough et al. 1979; Kammen et al. 1988; Wrzesinski et al. 1995). Therefore, the same tritium-labeled tRNA_Phe used in the TruB studies could also serve as a substrate for RluA and TruA. Time courses of pseudouridine formation were determined under single-turnover conditions by quench-flow measurements, as described for TruB. Again, 100% pseudouridine formation was observed after 5–10 sec for both enzymes (Fig. 4A,B). As before, the apparent rates were determined by one-exponential fitting and plotted against the enzyme concentration (Fig. 4C,D). Interestingly, the apparent rate of pseudouridine formation increased hyperbolically with increasing RluA concentration. This finding indicates that at low RluA concentrations substrate binding is limiting, which is overcome at higher RluA concentrations where catalysis is limiting. Therefore, we fit the concentration-dependence of the apparent rates, k_app, to a hyperbolic function in order to obtain the maximal rate at high RluA concentrations (k_max). This maximal rate corresponds to the rate constants of pseudouridine formation, k_p, and was determined to be 0.7 ± 0.15 sec⁻¹. For TruA, the apparent rate of pseudouridine formation, k_app, was not dependent on the enzyme concentration in the range tested, with an average rate of 0.35 ± 0.2 sec⁻¹ (Fig. 4D). This finding indicates that substrate binding is not rate-limiting for TruA under these experimental conditions, as previously observed for TruB. Hence, the apparent rate for pseudouridine formation by TruA reflects the rate constant of catalysis, k_p.

Uniform, slow catalysis of pseudouridine formation

The rate constants determined here for catalysis of pseudouridine formation, k_p, by RluA, TruA, and TruB are remarkably similar (Table 1). Given the precision of our measurements, the rate constants are almost identical and differ at most by a factor of two. The observation that catalysis is uniformly slow for all three pseudouridine synthases tested here raises the question whether catalysis is the rate-limiting step for these pseudouridine synthases. Since the apparent rates of pseudouridine formation reported here
for TruB and TruA are independent of the enzyme concentration (Figs. 3C, 4D), substrate binding is clearly not a limiting factor for TruB and TruA. Furthermore, the concentration dependence of pseudouridine formation by RluA (Fig. 4C) indicates that substrate binding is not limiting at concentrations above \( \sim 7.5 \) \( \mu \)M for this enzyme. This finding is in accordance with a previous report that binding is not the kinetically limiting step for yeast Pus1 (Arluison et al. 1999). The next question is whether product release could be limiting, which can be answered by comparing the elementary rate constant \( k_p \) and the catalytic constant \( k_{cat} \) determined under multiple-turnover conditions. The catalytic constant of TruA has been reported as 0.18 sec \(^{-1}\) (Huang et al. 1998), and under our experimental conditions an even higher value could be obtained by steady-state experiments (\( k_{cat} = 0.7 \pm 0.2 \) sec \(^{-1}\), data not shown). Thus, \( k_{cat} \) (0.18–0.7 sec \(^{-1}\)) and \( k_p \) (0.35 sec \(^{-1}\), vide supra) are comparable for TruA, indicating that product release is not overall rate-limiting, but that catalysis itself is the limiting step for TruA, as explained above for TruB. Thus, TruA and TruB resemble each other in this property. The catalytic constant \( k_{cat} \) for RluA has previously been measured to be \( \sim 0.1 \) sec \(^{-1}\) (Ramamurthy et al. 1999b), which is significantly lower than the rate constant for pseudouridine formation under single-turnover conditions determined here (\( k_p = 0.7 \) sec \(^{-1}\)). This indicates that, potentially, product release could be a rate-limiting step for RluA. It is not surprising that the enzymes might differ substantially in product release (and substrate binding) as they all display a different mode of specifically recognizing and interacting with their substrate RNA (Hoang and Ferré-D’Amaré 2001; Hoang et al. 2006; Hur and Stroud 2007). Therefore, it will

<table>
<thead>
<tr>
<th>Enzyme</th>
<th>( k_p ), sec (^{-1})</th>
</tr>
</thead>
<tbody>
<tr>
<td>RluA</td>
<td>0.7 ± 0.15</td>
</tr>
<tr>
<td>TruA</td>
<td>0.5 ± 0.2</td>
</tr>
<tr>
<td>TruB</td>
<td>0.35 ± 0.2</td>
</tr>
</tbody>
</table>

The rate constants for pseudouridine formation, \( k_p \), were determined from the enzyme concentration dependence of the apparent rates of single-turnover pseudouridine formation (Figs. 3C, 4C,D).
be interesting in the future to study the kinetic mechanism of RluA, TruA, and also TruB in greater detail using other stopped-flow techniques in order to analyze the mechanism of substrate recognition and product release by these pseudouridine synthases. Based on the data presented here, the mechanism of pseudouridine synthases consists of at least four steps: (1) substrate binding, (2) some conformational change such as tRNA unfolding and/or movement of the target uridine into the catalytic site, (3) catalysis consisting of several sub-steps (vide infra), and (4) product release. Further pre-steady-state kinetic analysis will reveal whether additional steps exist, as speculated previously (Arluison et al. 1999).

As the three pseudouridine synthases analyzed here are characterized by a uniformly slow rate constant for pseudouridylation (Table 1), it is conceivable that all pseudouridine synthases could have similar rate constants of \( \sim 0.5 \text{ sec}^{-1} \). All six families of pseudouridine synthases contain the same fold in the catalytic domain and very similar active sites including a catalytic aspartate (Hamma and Ferré-D’Amaré 2006). Therefore, it has been proposed that all pseudouridine synthases share a common catalytic mechanism (Hamma and Ferré-D’Amaré 2006); however, the catalytic mechanism is still not identified. Since RluA, TruA, and TruB represent three different families of pseudouridine synthases, it is rather likely that also the other enzyme families share a rather low rate constant for pseudouridylation. Interestingly, a catalytic rate constant of \( \sim 0.5 \text{ sec}^{-1} \) is rather small compared to many other enzymes which often achieve \( k_{\text{cat}} \) values of \( 10^2–10^6 \text{ sec}^{-1} \) (Voet and Voet 2011). The rate enhancements by pseudouridine synthases cannot be quantitatively determined since no data exist on the rate of the uncatalyzed reaction, maybe because this reaction would not occur without catalysis. Assuming that the uncatalyzed reaction is very slow or not occurring at all, the rate enhancement by pseudouridine synthases might be significant despite the relatively low catalytic rate constant \( k_{\text{q}} \).

Three reasons can be envisioned to explain the low rate constant of pseudouridylation. First, the low rate constant of pseudouridylation might be due to the absence of evolutionary pressure to further increase the rate of pseudouridylation. While pseudouridines are the most common RNA modifications and supposed to enhance RNA structure and function, many pseudouridines and, in turn, many pseudouridine synthases, are not essential for the cell (Raychaudhuri et al. 1999; Gutgsell et al. 2000; Del Campo et al. 2001; Kinghorn et al. 2002). Second, it might even be envisioned that pseudouridine synthases have been selected to be slow. Such a selection could arise if another function in addition to pseudouridylation is important for this enzyme family, such as a role in RNA folding which has been suggested previously (Hoang and Ferré-D’Amaré 2001). Third, it might not be possible to further increase the rate constant of catalysis of pseudouridylation due to the actual chemistry of the reaction. Pseudouridine synthases catalyze a challenging chemical reaction consisting of multiple steps including cleavage of the N1-glycosidic bond, rotation of the uracil base, and formation of a new C5-glycosidic bond. To the best of our knowledge, this complex isomerization of uridine to pseudouridine is irreversible. It is remarkable that these enzymes efficiently catalyze these three different chemical reactions in the same catalytic pocket. These restrictions might impose an upper limit on the achievable rate constant for catalysis. In this case, all pseudouridine synthases would be limited by the same chemical difficulty and most likely display the same rate constants for catalysis—as observed here for RluA, TruA, and TruB. More investigations are necessary to distinguish among the three chemical reactions in pseudouridine catalysis and to identify the rate-limiting step within these reactions. Importantly, the quench-flow technique used here might help to isolate and characterize transient intermediates on the reaction pathway (Barman et al. 2006). Interestingly, the first substep, cleavage of the N-glycosidic bond, resembles the reaction catalyzed by uracil-DNA glycosylases, which display \( k_{\text{cat}} \) values of \( 4–200 \text{ sec}^{-1} \) (Duraffour et al. 2007; Liu et al. 2007), at least 10-fold higher than the catalytic rate constant of pseudouridine formation reported here. Therefore, N-glycosidic bond cleavage can be fast in principle; and the subsequent steps of base rotation or C-C bond formation are more likely the limiting steps during pseudouridylation.

In conclusion, we present here the first pre-steady-state rapid kinetic analysis of pseudouridine synthases. Thereby, important insight into the kinetic mechanism of TruB has been obtained revealing a two-step substrate binding and slow, rate-limiting catalysis of pseudouridylation. This two-step binding mechanism might be common among RNA modification enzymes and might contribute to the specific recognition of selected target sites. Furthermore, our kinetic analysis of RluA, TruA, and TruB representing three different families of pseudouridine synthases demonstrated that catalysis of pseudouridine formation is a uniformly slow step (Table 1), which is most likely a general feature of all pseudouridine synthases. These findings are pivotal for the analysis and dissection of the catalytic mechanism and the kinetics of the three chemical substeps taking place during pseudouridylation. Moreover, it will be interesting to compare the presented kinetic mechanism of bacterial, stand-alone pseudouridine synthases with eukaryotic homologs, as well as with H/ACA small ribonucleoproteins, which employ a different approach to substrate RNA binding by base-pairing to an H/ACA guide RNA and might, therefore, have a different rate-limiting step such as product release (Li 2008). In summary, the presented pre-steady-state analysis of the basic kinetic mechanism of pseudouridylation identified for the first time catalysis as a slow step in three pseudouridine synthase families and lays the groundwork for future investigations on the detailed kinetic mechanism of H/ACA small ribonucleoproteins, on the mechanism of substrate binding by TruB and other stand-alone pseudouri-
dine synthases, and on the catalytic mechanism of pseudouridine formation in general.

MATERIALS AND METHODS

Buffers and reagents

Buffer TAKEM4: 50 mM Tris-HCl pH 7.5, 70 mM NH₄Cl, 30 mM KCl, 1 mM EDTA, 4 mM MgCl₂, Nucleotide triphosphates and guanine monophosphate for in vitro transcription, DNasel, and inorganic pyrophosphatase were from Sigma; all other enzymes were from Fermentas. Chemicals were purchased from VWR. DNA oligos were obtained from IDT, and radioactive UTP was from Moraveck.

Molecular cloning and mutagenesis

The open reading frame of E. coli truB was ligated into pET28(+) vector encoding an N-terminal histidine-tag using restriction sites Nhel and BamHI to generate the plasmid pET28a-truB (similar to Nurse et al. [1995]). The QuickChange method (Strategene) was used for site-directed mutagenesis generating plasmid pET28a- TruBD48N. Gene sequences were confirmed by DNA sequencing (Macrogen).

Protein expression and purification

For protein expression, pET28a-TruB and pET28a-TruBD48N were transformed into BL21(DE3) competent E. coli cells (EMD Bioscience) which were grown in LB medium supplemented with 50 μg/mL kanamycin at 37°C. TruA and RluA were expressed using the plasmids pCA24N(GFP minus)-JW0057 in AG1(ME5305) plasmids pCA24N(GFP minus)-KCl, 1 mM EDTA, 4 mM MgCl₂. Nucleotide triphosphates and guanine monophosphate for in vitro transcription, DNasel, and inorganic pyrophosphatase were from Sigma; all other enzymes were from Fermentas. Chemicals were purchased from VWR. DNA oligos were obtained from IDT, and radioactive UTP was from Moraveck.

In vitro transcription

The template for the in vitro transcription of E. coli tRNA^phc was generated by PCR amplification from the plasmid pCFO (Sampson et al. 1989) (kind gift of O. Uhlenbeck) using the following primers: 5’-GCTGCAGTAATACGACTCACTATAG-3’ and 5’-mUmGTTGCCGGACTCG-3’.

Subsequently, the in vitro transcriptions were performed using the PCR template [10% (v/v)] in transcription buffer (40 mM Tris-HCl pH ~7.5, 15 mM MgCl₂, 2 mM spermidine, 10 mM NaCl, 10 mM DTT) with 3 mM NTPs (ATP, CTP, GTP, and UTP), 5 mM GMP, 0.01 U/μL inorganic pyrophosphatase, 0.3 μM T7 RNA Polymerase, and 0.12 U/μL RNase inhibitor at 37°C for 16 h. For generation of [3H]-labeled tRNA, the in vitro transcriptions were performed for 4 h using 3 mM ATP, CTP, and GTP each, and 0.1 mM [5-3H]UTP (0.46 Ci/m mole). Following the in vitro transcription step, the template was digested with DNasel for 1 h at 37°C. The nonradioactive RNA was purified by DEAE anion exchange chromatography (Easton et al. 2010). [3H]-labeled RNA was purified with a Nucleobond AX20 column (Macherey-Nagel) using equilibration buffer R0 [100 mM Tris-acetate pH 6.3, 10 mM MgCl₂, 15% (v/v) ethanol], washing buffer R1 (R0 plus 300 mM KCl), and elution buffer R3 (R0 plus 1150 mM KCl). The obtained tRNA was concentrated by isopropanol precipitation and dissolved in H₂O. The tRNA concentration was determined photometrically at 260 nm using the extinction coefficient 5 \times 10^5 M⁻¹ cm⁻¹ (Peterson and Uhlenbeck 1992). The specific activity of the purified tRNA was determined by scintillation counting.

Nitrocellulose filtration

Prior to the experiment, folding of tRNA^phc was allowed to occur by heating 2 μM [3H]tRNA^phc in TAKEM₄ buffer to 60°C for 5 min and subsequent slow cooling to 37°C (Hengesbach et al. 2010). To allow for tRNA binding to TruB D48N, 50 nM [3H]tRNA^phc were incubated with 0–30 μM TruB D48N in TAKEM₄ buffer for 10 min at room temperature. The complete 50 μL reaction mixture was then filtered through a nitrocellulose membrane followed by washing of the membrane with 1 mL cold TAKEM₄ buffer. Membranes were dissolved for 30 min in 10 mL EcoLite scintillation cocktail [EcoLite (+), MP Biomedical], and the amount of tRNA bound to TruB D48N retained on the membrane was determined by scintillation counting (Perkin-Elmer Tri-Carb 2800TR liquid scintillation analyzer). In order to obtain the dissociation constant (Kₐ), the increase in the fraction of bound tRNA as a function of the TruB D48N concentration was analyzed by fitting to a hyperbolic equation:

\[
\text{Bound} = \text{Bound}_{\text{max}} \times \left( \frac{\text{TruBD48N}}{[\text{KD} + \text{TruBD48N}]} \right)
\]

Absorbance spectroscopy and stopped-flow measurements

Following folding of the tRNA as described above, the absorbance at 260 nm of a 0.8 μM tRNA solution in TAKEM₄ was recorded.
Increasing amounts of TruB D48N were added to the tRNA solution which was incubated at room temperature for 1 min, and the absorbance increase at 260 nm was monitored. The same titration of TruB D48N into TAKEM4 buffer was performed, and the resulting absorbance readings were subtracted from the data in the presence of tRNA, yielding the increase in absorbance at 260 nm due to the interaction of tRNA$^{\text{Phe}}$ and TruB D48N. This change in absorbance was then plotted against the TruB D48N concentration, and the data were subsequently analyzed by fitting with a quadratic function \((\Delta A_{260} = A_0 + \text{Amp} \times [\text{RNA}] / [\text{TruB}]/2 - (K_D + [\text{RNA}] / [\text{TruB}])^2 / 4 - [\text{TruB}] \times [\text{RNA}]^{0.5})\)

A quadratic function was used for fitting instead of a hyperbolic function since the RNA concentration was not significantly lower than the TruB concentration used.

Pre-steady-state kinetics of tRNA interaction with TruB were monitored in a KinTek SF-2004 stopped-flow apparatus. Twenty five μL folded tRNA$^{\text{Phe}}$ (final concentration 0.75 μM) were rapidly mixed with 25 μL TruB (final concentration 2.5–10 μM) at 20°C in TAKEM4, and the absorbance at 260 nm was recorded. The starting absorbance was subtracted, and the resulting time courses were analyzed by fitting with a one-exponential function to determine the apparent rate \(k_{app}\):

\[
A = A_0 + \text{Amp} \times \exp(-k_{app} \times t)
\]

The apparent rates \(k_{app}\) were then plotted against the enzyme concentration.

**Tritium release assay to detect pseudouridylation**

Prior to all experiments, \(\text{[3H]}\)tRNA$^{\text{Phe}}$ was allowed to fold as described above. For Michaelis-Menten experiments, different concentrations of \(\text{[3H]}\)tRNA$^{\text{Phe}}$ (100–2000 nM) were incubated with 10 nM enzyme in TAKEM4 buffer at 37°C, and samples (10.8 pmol \(\text{[3H]}\)tRNA$^{\text{Phe}}$) were removed after 30 sec, 60 sec, and 120 sec. The reaction was stopped by adding the samples to 1 mL 5% (w/v) activated charcoal (Norit A) in 0.1 M HCl. Following centrifugation at 10,000 X g for 2 min, 0.8 mL of the supernatant was added to 0.5 mL fresh 5% Norit A (w/v) in 0.1 M HCl, mixed, and centrifuged again. One mL of the supernatant was filtered through glass wool plugged in a 1-mL micropipet tip, and 0.8 mL of the resulting filtrate was then used for scintillation counting in 4 mL EcoLite scintillation cocktail. The concentration of released tritium corresponding to the formed pseudouridine was calculated and divided by the respective incubation time, yielding the initial rate of the reaction. The dependence of the initial rates \(v_0\) on the tRNA concentration was fitted with a Michaelis-Menten equation

\[
v_0 = v_{max} \times [\text{tRNA}^{\text{Phe}}] / (K_M + [\text{tRNA}^{\text{Phe}}])
\]

with \(v_{max} = k_{cat} \times [\text{enzyme}]\).

Pre-steady-state measurements were performed in a KinTek quench-flow apparatus by rapidly mixing 14.5 μL folded \(\text{[3H]}\)-tRNA$^{\text{Phe}}$ (final concentration 1.0 μM) with 13 μL of enzyme (final concentration 2.5–15 μM) at 37°C in TAKEM4. The reaction was stopped at desired time points by 0.1 M HCl. The total \(\text{[3H]}\)tRNA$^{\text{Phe}}$ concentration was determined by liquid scintillation counting of 2 μL of the quenched sample. To measure the concentration of released tritium, a defined volume (120–220 μL) of the quenched sample was added to 1 mL 5% Norit A (w/v) in 0.1 M HCl, and processed as described above. The percentage of pseudouridine formation was calculated from the total \(\text{[3H]}\)tRNA$^{\text{Phe}}$ concentration and the concentration of released tritium for each time point. Fitting of the time courses with a one-exponential equation

\[
F = F_0 + A \times \exp(-k_{app} \times t)
\]

yielded the apparent rate, \(k_{app}\), of pseudouridine formation, which then was plotted against the enzyme concentration. For RluA, this concentration dependence was fit to a hyperbolic equation to obtain the maximal rate \(k_{max}\):

\[
k_{app} = k_{max} \times [\text{RluA}] / (K_{half} + [\text{RluA}])
\]

**ACKNOWLEDGMENTS**

We thank Laura Hagstrom and Theron White for their help with cloning and initial purification of TruB; Olke Uhlenbeck (Northwestern University, Evanston, IL) for plasmid pCFO; the National BioResource Project (NIG, Japan) for the TruA and RluA expression plasmids; and Hans-Joachim Wieden for providing access to the quench-flow and stopped-flow apparatus as well as for critically reading the manuscript. This work was supported by the National Science and Engineering Research Council of Canada (NSERC), and the Canada Foundation for Innovation (CFI).

Received July 3, 2011; accepted August 29, 2011.

**REFERENCES**


Pre-steady-state kinetic analysis of the three *Escherichia coli* pseudouridine synthases TruB, TruA, and RluA reveals uniformly slow catalysis

Jaden R. Wright, Laura C. Keffer-Wilkes, Selina R. Dobing, et al.

*RNA* 2011 17: 2074-2084 originally published online October 13, 2011
Access the most recent version at doi:10.1261/rna.2905811