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Vinyluridine as a Versatile Chemoselective Handle for the Posttranscriptional Chemical Functionalization of RNA

Jerrin Thomas George and Seergazhi G. Srivatsan*

Department of Chemistry, Indian Institute of Science Education and Research (IISER) Pune,
Dr. Homi Bhabha Road, Pashan, Pune 411008, India.

ABSTRACT: Development of modular and efficient methods to functionalize RNA with biophysical probes is very important to advance the understanding of the structural and functional relevance of RNA in various cellular events. Herein we demonstrate a two-step bioorthogonal chemical functionalization approach for the conjugation of multiple probes onto RNA transcripts using a 5-vinyl-modified uridine nucleotide analog (VUTP). VUTP, containing a structurally non-invasive and versatile chemoselective handle, was efficiently incorporated into RNA transcripts by *in vitro* transcription reactions. Further, we show for the first time the use of a palladium-mediated oxidative Heck reaction in functionalizing RNA with fluorogenic probes by reacting vinyl-labeled RNA transcripts with appropriate boronic acid substrates. The vinyl label also permitted the posttranscriptional functionalization of RNA by a reagent-free inverse electron demand Diels-Alder (IEDDA) reaction in the presence of tetrazine substrates. Collectively, our results demonstrate that the incorporation of VUTP provides newer possibilities for the modular functionalization of RNA with variety of reporters.

INTRODUCTION

Functional modification of RNA *in vitro* and in cells have become necessary for understanding the regulation, structure, and function of RNA.¹⁻³ While solid-phase oligonucleotide (ON) synthesis and enzymatic labeling methods using bacteriophage RNA polymerases are widely used in the generation of labeled RNA suitable for cell-free and in-cell analysis, analogous approaches to label endogenous RNA are less prevalent.⁴ Immunostaining of metabolically incorporated nucleoside analogs (e.g., BrU) using antibodies, structure-specific antibodies (e.g., G-quadruplexes), aptamers (e.g., spinach) and dyes have been employed to visualize cellular RNA structure and synthesis.⁵⁻¹⁰ However, limited permeability and selectivity of the antibodies and compromised function of aptamer-tagged RNA have been the downsides of these methods. Alternatively, bioorthogonal

chemical reactions provide an easy route to label RNA *in vitro* and cellular milieu for a variety of applications.¹¹⁻¹⁵ In this approach, a reactive group is introduced into an RNA ON by either chemical, enzymatic (RNA polymerases) or chemo-enzymatic (transferases) means and further bioconjugation to label RNA is achieved by performing a chemoselective reaction with a cognate reactive partner containing a desired biophysical reporter. Several chemoselective reactions including azide-alkyne cycloaddition (AAC), Staudinger ligation, and IEDDA reactions have been effectively utilized in labeling protein, glycan, lipid and DNA.¹⁶⁻¹⁸ However, establishing RNA labeling techniques under the conditions used in these reactions has always been a challenge due to the inherent instability of the RNA.¹⁵

Among the various bioorthogonal reactions, AAC reaction has been extensively used in labeling RNA postsynthetically.¹⁹⁻²⁷ Typically in this reaction, a copper (I) stabilizing ligand is used to alleviate the toxic effects of copper ions, which is known to produce species harmful to nucleic acids in the redox environment.^{14,28,29} In another strategy, the use of toxic copper has been circumvented by employing strain-promoted AAC (SPAAC) reaction, wherein strained alkynes (e.g., cyclooctynes) are used as one of the reactive counterparts.³⁰⁻³⁴ However, SPAAC reaction efficiency in labeling nucleic acids in cells has been limited by the poor permeability and reactivity of bulky cyclooctyne probes.^{33,34} Other concerns which have limited the application of AAC reactions are the involvement of alkyne substrates in side reactions such as homocoupling and thiol-yne addition, and low stability of azide substrates in the chemical labeling conditions.^{33,35,36}

Analogous to azide and alkyne groups, the ability of alkene group to participate in a wide range of chemoselective reactions has been elegantly utilized in devising several bioconjugation strategies. Bioconjugation methods based on ene-thiol and ene-tetrazole reactions have been put to good use in cross-linking and functionalizing biomolecules, especially proteins.³⁷⁻³⁹ However, the use of UV radiation and non-specific reaction with cellular thiols has hampered their applications in in-cell analysis. Recently, protein labeling has been achieved by using oxidative Heck reaction between protein-bound alkenes and boronic acid reporters.⁴⁰⁻⁴² This reaction is particularly interesting because it can be effectively performed in aqueous buffer under near physiological conditions (~pH 7, room temperature and oxygen atmosphere) as compared to other analogous Pd-assisted biomolecular labeling reactions.^{43,44} Very recently, base-modified 2'-deoxynucleosides, containing a minimally perturbing vinyl group as a dienophile, have been successfully incorporated *in vitro* into DNA ONs and metabolically into replicating DNA in cells.^{45,46}

Although the reaction rates are significantly lower as compared to strained dienophiles, vinyl-labeled DNA did enable the labeling as well as visualization of DNA in cell-free and cellular environments by using IEDDA reaction.

Coupling reactions have been employed to attach heterocyclic aryl moieties via an alkene to natural nucleobases. Such conjugations have produced microenvironment-sensitive nucleoside probes, which have been used in studying the conformation of the G-quadruplex structure.⁴⁷ Similarly, fluorogenic reporters have been developed to specifically detect epigenetic and epitranscriptomic nucleobase modifications (e.g., 5-formylcytosine and 5-formyluracil) by using a chemoselective reaction between the base and trimethylindole derivatives.⁴⁸ Taking advantage of ~~these~~ this knowledge, we rationalized that enzymatic incorporation of a minimally perturbing vinyl-modified ribonucleotide into RNA would facilitate the generation of microenvironment-sensitive fluorogenic RNA reporters by posttranscriptional oxidative Heck reaction with appropriate boronic acid derivatives (Figure 1). Also, we envisioned that vinyl-labeled RNA transcript would open up possibilities to investigate its potential to functionalize RNA by reagentless IEDDA reaction. Despite the successes with protein and DNA,^{40,49} the utility of vinyl group as a versatile chemoselective handle in developing RNA labeling protocols has not been well explored.⁵⁰ In this context, here we describe the synthesis of 5-vinyl-modified UTP analog (VUTP) and its effective incorporation into RNA ONs by *in vitro* transcription reactions. Further, we illustrate the posttranscriptional chemical functionalization of vinyl-labeled RNA transcripts with a variety of biophysical reporters by using oxidative Heck and IEDDA reactions. Notably, this is the first report on the bioconjugation of an RNA ON by employing Pd-mediated oxidative Heck reaction.

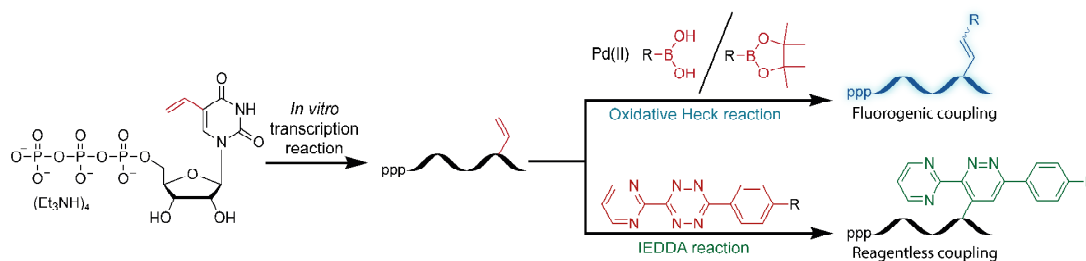
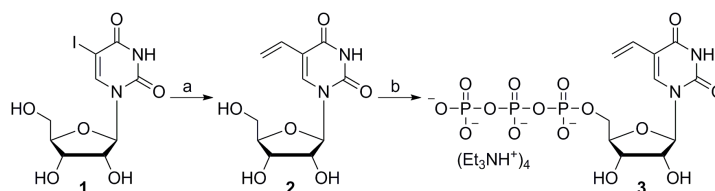


Figure 1. Diagram illustrating the posttranscriptional chemical labeling of RNA by using oxidative Heck and IEDDA reactions. Incorporation of VUTP into RNA transcripts, followed by oxidative Heck reaction with boronic acid derivatives and IEDDA reaction with tetrazole derivatives enabled the synthesis of RNA conjugated to various reporters/tags.

RESULTS AND DISCUSSION

Synthesis and enzymatic incorporation of 5-vinyluridine triphosphate (VUTP) 3. VUTP **3** was synthesized in simple steps starting from 5-iodouridine **1** (Scheme 1). 5-Vinyluridine (VU) **2** was synthesized by Stille cross-coupling reaction using vinyltributylstannane and a palladium catalyst, tris(dibenzylideneacetone)dipalladium(0).⁵¹ VU was further phosphorylated in the presence of POCl₃ and *bis*-tributylammonium pyrophosphate to afford VUTP. The ability of VUTP to serve as a UTP analog for RNA polymerase was studied by performing *in vitro* transcription reactions in the presence of a small series of DNA templates **S1–S4** (Figure 2). The templates contained one or two dA residues in the coding region to guide the insertion of monophosphate of VUTP into RNA during polymerization reaction. The 5'-end of each template ended with a dT residue to introduce a radiolabeled α -³²P adenosine at the 3'-end of the full-length transcript. Transcription was performed using GTP, CTP, UTP/**3**, α -³²P ATP in the presence of T7 RNA polymerase. The transcription products were then electrophoresed on an analytical polyacrylamide gel and phosphor imaged.



Scheme 1. Synthesis of VUTP **3**. (a) Pd₂(dba)₃, P(Fur)₃, vinyltributylstannane, DMF, 60 °C, 3 h, 53 %. (b) (i) POCl₃, proton sponge, trimethylphosphate, ~4 °C, 30 min, (ii) *bis*-tributylammonium pyrophosphate, tributylamine, ~4 °C, 30 min, 21 %.

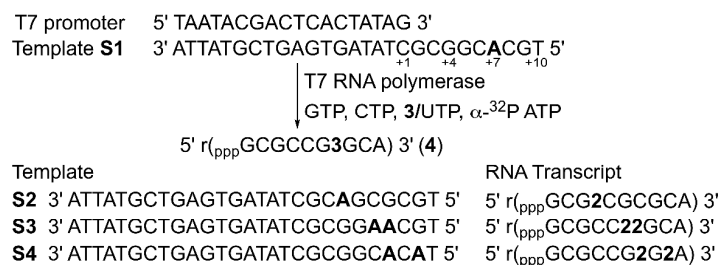


Figure 2. Incorporation of vinyl group into RNA ONs by *in vitro* transcription reaction using DNA templates **S1–S4** and VUTP **3**. “r” preceding the sequence in parenthesis represents RNA ON transcripts.

The incorporation of VUTP **3** in the presence of template **S1** proceeded with very good efficiency at +7 position yielding a 10 mer full-length RNA transcript **4** (Figure 3, lane 2). Slower migration of the transcription product **4** in lane 2 indicated hinted the

incorporation of **3** into the transcript (compare lane 1 and 2). While a control reaction in the absence of UTP and **3** did not yield full-length RNA transcript, at equimolar concentrations, both natural UTP and **3** got incorporated into RNA (Figure 3, lane 3 and 4, respectively). Templates **S2–S4** incorporated modified UTP near the promoter region and at more than one site with good to excellent efficiency (Figure 3, lane 6, 8 and 10).

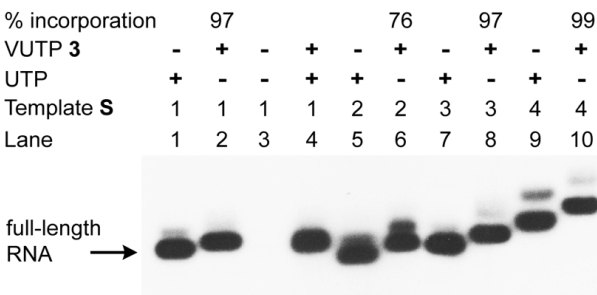


Figure 3. Phosphor image of transcripts obtained by transcription of templates **S1–S4** in the presence of UTP and or **3**. % Incorporation of **3** is reported relative to a control transcription with UTP. For complete gel picture see Figure S1.

Transcript **4** was scaled up by performing large-scale reactions with **S1** in the presence of cold NTPs. Typically, a reaction in the presence of 75 pmol of the template gave nearly 15 nmol of the modified transcript after gel electrophoretic purification, which was further used for posttranscriptional chemical labeling experiments. The presence of VU in transcript **4** was confirmed by mass analysis and enzymatic digestion assay (Figure S2, S3 and Table S1).

Oxidative Heck reaction on vinyl-labeled RNA transcript. Electronically unbiased terminal alkene (e.g., allylic) substrates have been reported to undergo oxidative Heck reaction with boronic acids in the presence of water-soluble Pd-EDTA complex.⁴¹ But the feasibility of such a reaction on RNA has not been explored. In this regard, we sought to investigate the possibility of performing oxidative Heck reaction on vinyl-modified RNA with suitable boronic acid and ester substrates. For this purpose we chose a range of heterobicycles, namely benzothiophene-2- (**5**), benzofuran-2- (**6**), indole-5- (**7**), dibenzothiophene-4-boronic acid (**8**) and a previously reported pinacol boronic ester, benzothiophene-2-vinyl boronic ester (**9**),⁴⁷ which upon conjugation to uracil ring could impart fluorescence to otherwise non-emissive nucleobase (Figure 4). Fluorescent nucleosides have been developed by attaching heterocycles to pyrimidine and purine bases,

respectively, and such analogs incorporated into ONs have been used as probes in various nucleic acid studies.^{2,52-57}

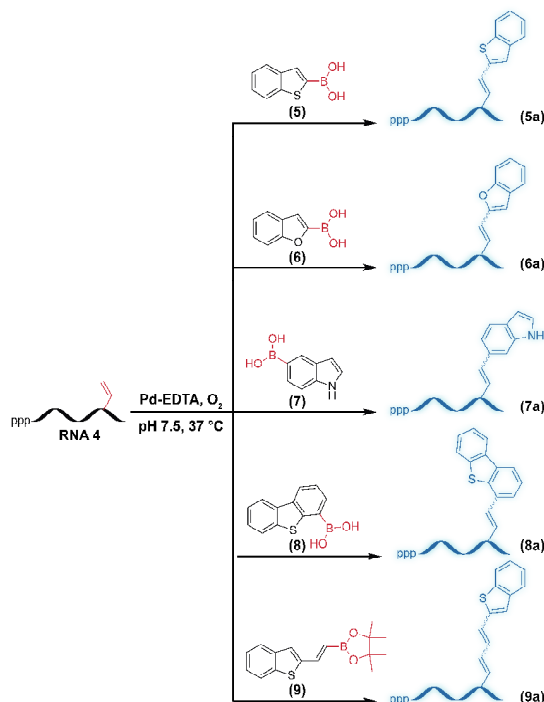


Figure 4. Oxidative Heck reaction between vinyl-modified RNA ON **4** and boronic acid substrates **5–8** or boronic ester **9** in the presence of Pd-EDTA complex.

For performing oxidative Heck reaction on RNA, transcript **4** was incubated with boronic acid **5** in the presence of Pd-EDTA complex at 37 °C. Aliquots of reaction mixture at various time intervals were resolved by PAGE under denaturing conditions. Almost complete conversion was observed at 18 h, and rewardingly, we observed an intense fluorescent band corresponding to the coupled product along with a minor band (Figure 5 and Figure S4). Reactions with other substrates (**6–8**) also proceeded well (Figure S5). Further, products from large-scale reactions between transcript **4** and boronic acid **5–8** or boronic ester **9** were analyzed by RP-HPLC at 260 nm and 338 nm (conjugation of heterocycles to pyrimidine bases results in a strong absorption band near 338 nm).^{2,58} Mass analysis of HPLC fractions corresponding to both 260 nm and 338 nm absorption bands revealed the formation of the oxidative Heck products **5a–9a** (Figure S6, S7 and Table S2). Reactions performed at a 5 nmole scale of the RNA ON **4** provided 0.5 to 1.1 nmole of the coupled products (Table S3).

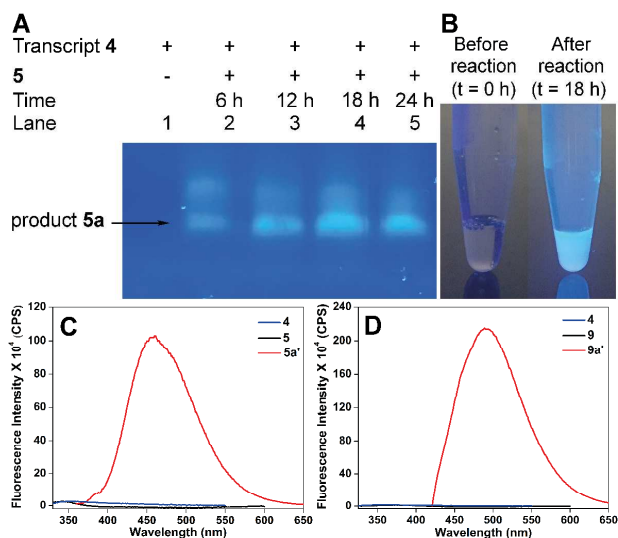


Figure 5. (A) UV-transilluminator image (364 nm) of the gel of a reaction between RNA 4 and 5. See Figure S4 for the UV-shadow image of the gel at 254 nm. (B) Fluorescence of RNA transcript before and after the reaction between RNA 4 and boronic acid 5 under UV lamp (364 nm). (C) Fluorescence spectra (0.2 μ M) of RNA transcript 4, boronic acid 5 and RNA product 5a (HPLC fraction corresponding to 5a'). Samples were excited at 289 nm, 310 nm and 340 nm, respectively, with an excitation and emission slit width of 8 nm and 9 nm. (D) Fluorescence spectra (0.2 μ M) of RNA transcript 4, boronic ester 9 and RNA product 9a (HPLC fraction corresponding to 9a'). Samples were excited at 289 nm, 310 nm and 372 nm, respectively, with an excitation and emission slit width of 7 nm and 8 nm. See Figure S6 for HPLC profiles.

An oxidative Heck reaction between an electronically unbiased terminal alkene (e.g., allylic system) and arylboronic acid substrates, typically gives linear and branched cross-coupled products.⁴⁰ However, reactions with electronically biased aryl vinyl systems (e.g., styrene) give predominantly linear trans-coupled product.^{59,60} Formation of branched regioisomer product has also been reported using Pd in the presence of certain bulky ligands (e.g., 2,9-dimethyl-1,10-phenanthroline).⁶⁰ Although it is not a major concern in most bioconjugation strategies, we sought to determine the formation of different isomers in the oxidative Heck reaction of vinyl-labeled RNA transcript. First, we carried out a reaction between free vinyl-modified uridine 2 and boronic acid 5 under similar conditions used for RNA ligation. Interestingly, at the ribonucleoside level a very low conversion was observed even after prolonged incubation time. Varying the Pd-ligand complex and oxidant did not give a better yield (data not shown). However, a fluorescent product was isolated in analytical quantities, and further characterization by NMR revealed the formation of a linear trans-isomer product 2a (Figure S8). Further, the coupled product gave a unique longer wavelength absorption band at \sim 338 nm as compared to native nucleoside/nucleic acid, which absorbs at 260 nm (Figure S8B). Next, the major HPLC fraction corresponding to the

RNA products (designated as **5a'**) of a reaction between vinyl-labeled RNA ON **4** and boronic acid **5** was subjected to enzymatic digestion in the presence of calf intestinal alkaline phosphatase, RNase A, RNase T1 and snake venom phosphodiesterase. Enzymatic digestion of **5a'** gave individual ribonucleosides as well as Heck-coupled ribonucleosides, whose identity was determined by comparing the HPLC chromatogram of the digested sample and oxidative Heck reaction product **2a** obtained from a reaction between uridine **2** and boronic acid **5** (Figure S9). The chromatogram of the digest revealed the presence of native ribonucleosides (rC, rG and rA) and two peaks corresponding to coupled products absorbing at 338 nm. One of the peaks clearly matched with the trans-isomer product **2a**, whereas the chemical structure of the peak **X** could not be determined. However, both **2a** and **X** gave the same mass indicating that that a mixture of isomers (**X** may be cis isomer) are formed in these reactions. Formation of a mixture of ~~regio—and-stereoisomers~~ isomers, which is an inherent “limitation” of this reaction, has also been observed in protein ligation experiments.^{40,41} Although, at the present, this reaction may not be viable to introduce a specific type of conformation-sensitive probe into RNA, development in chelate-control strategies could vastly improve the selective of the reaction.⁶⁰

We next studied the fluorogenic nature of the RNA products by steady-state fluorescence spectroscopy. Benzothiophene-coupled RNA product **5a** (HPLC fraction corresponding to **5a'**) showed a ~40-fold enhancement in fluorescence intensity as compared to the boronic acid substrate **5** (Figure 4, 5C and Figure S6). Interestingly, benzothiophene-alkene-coupled RNA product **9a** (HPLC fraction corresponding to **9a'**) showed a remarkable ~170-fold enhancement in fluorescence intensity as compared to substrate **9**. Further, the emission profile of **9a** showed a significant bathochromic shift compared to RNA **5a** as a direct result of increased conjugation between the uracil and benzothiophene rings (Figure 4, 5D and Figure S6). Reactions with benzofuran (**6**), indole (**7**), dibenzothiophene (**8**) boronic acid substrates also produced fluorescent RNA products (Figure S10). While benzofuran- and dibenzothiophene-conjugated RNA products were reasonably fluorescent, indole-conjugated RNA was found to be weakly fluorescent. Taken together, this approach of functionalizing vinyl-labeled RNA transcripts by fluorogenic Heck-type coupling reaction with appropriate boronic acid/boronic ester substrates could provide direct access to RNA functionalized with fluorescent reporters.^{48,61, 62}

IEDDA reaction on vinyl-labeled RNA transcripts. Analogous to AAC chemistry, IEDDA reaction between an electron-rich dienophile and electron-deficient tetrazine is gaining particular attention as a useful biomacromolecular labeling strategy because this reaction is reagent free, reasonably fast, highly selective and biocompatible.⁶³⁻⁶⁵ Bioconjugation strategies based on IEDDA reaction commonly use ONs labeled with reactive dienophiles such as norbornene, trans-cyclooctene and cyclopropene groups.⁶⁶⁻⁷³ While reaction rates of strained dienophiles with tetrazine derivatives are generally fast, such bulky dienophile substituents on nucleoside are particularly not suitable for labeling cellular nucleic acids as they may not serve as good substrates for endogenous polymerases. Albeit low reactivity as compared to strained dienophiles, vinylated 2'-deoxynucleosides have been used as chemical reporters to label and visualize DNA in cells by IEDDA reaction.⁴⁵ Encouraged by these results, we decided to study the reactivity of vinyl label of transcript **4** towards diene counterpart in IEDDA reaction. For this purpose, we chose two tetrazine cores, which had reportedly shown a good reactivity against an electron-rich strained alkene.⁶⁶⁻⁷³ The reactivity of transcript **4** was tested by using a commercially available tetrazine **10**, and biotinylated (**11**) and Cy5-conjugated (**12**) tetrazines, which were synthesized by following an analogous literature procedure (Figure 6 and Scheme S1).⁶⁴

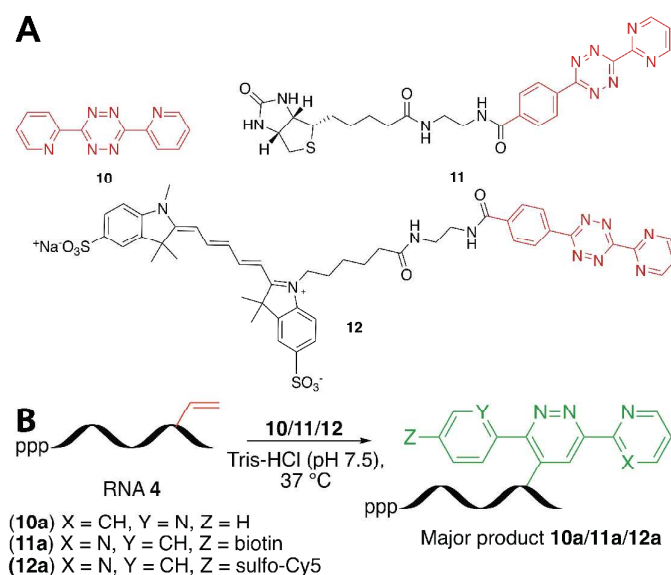


Figure 6. (A) Tetrazine substrates **10–12** used in this study. (B) IEDDA reaction between RNA ON **4** and tetrazine substrates. **10a–12a** are RNA products obtained from a reaction between RNA ON **4** and tetrazines **10–12**, respectively.

Initially, the IEDDA reaction was performed using the free nucleoside **2** and a tetrazine substrate **10**. The reaction proceeded well and almost complete conversion was observed after 21 h (Scheme S2 and Figure S11). The reaction resulted in a mixture of ligated products (**2b** and **2c**), which were isolated and characterized. **2a** was found to be oxidized pyridazine product, whereas **2c** corresponded to the dihydropyridazine product, which underwent slow oxidation to form **2b**. Formation of a mixture of products and slow oxidation event has been documented earlier in DNA conjugation experiments.^{45,74}

Next, IEDDA reactions were performed between RNA ON **4** and tetrazine substrates in Tris-HCl buffer (pH 7.5) at 37 °C, and aliquots of reaction mixture at various time intervals were analyzed by analytical PAGE under denaturing conditions. A reaction with tetrazine **10** was found to be almost complete in 12 h, whereas with substrates **11** and **12** the reaction was partially complete even after 15 h (Figure S12). Reactions at elevated temperatures (45 °C and 55 °C) did not result in noticeable improvement in reaction efficiency (data not shown). This is in agreement with reports of less reactivity of the tetrazine core of **11** and **12** as compared to tetrazine core of **10**.⁶⁵

Large-scale reactions were then performed to isolate and characterize the RNA products. Reaction with substrates **10–12** produced major and minor (could not be isolated) bands (Figure S13). Mass analysis of major bands confirmed the formation of pyridazine products **10a–12a** (Figure S14 and Table S2) similar to previous report.⁷² UV-vis profile of purified products was distinguishingly different from the profile of transcript **4**, and further fluorescence analysis also confirmed the bioconjugation by IEDDA reaction (Figure S15). While HPLC analysis of the major band obtained from a reaction with symmetrical tetrazine **10** gave a single peak corresponding to the pyridazine **10a**, reactions with asymmetrical tetrazines **11** and **12** afforded mixture of pyridazine isomers (Figure S16). The formation of mixture of isomeric products is consistent with literature reports.^{45,75}

High density labeling of VUTP into a longer RNA transcript was studied by using a DNA template that would generate a 59 mer RNA containing 9 modifications. T7 RNA polymerase effectively incorporated the vinyl analog into the 59 mer RNA **13** with efficiency comparable to that of natural UTP (Figure 7A). Transcript **13** was subsequently reacted with sulfo-Cy5 labeled tetrazine **12** and the reaction product was resolved on a 2% agarose gel. The image captured using UV-transilluminator (364 nm) clearly revealed the fluorescence labeling of the longer RNA transcript (Figure 7B).

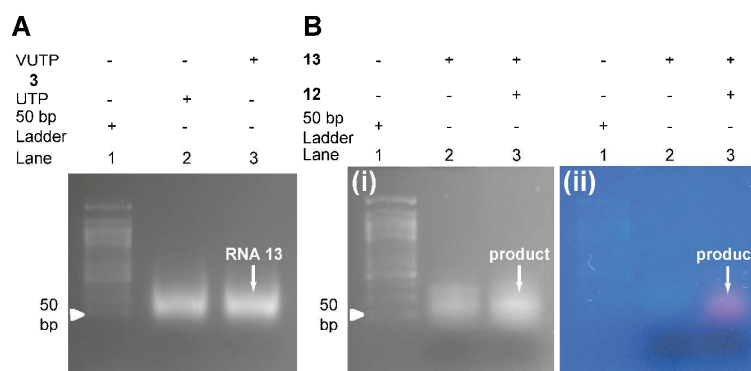


Figure 7. (A) Agarose gel picture of 59 mer control (lane 2) and VU-modified RNA transcript **13** (lane 3). Lane 1: 50 bp DNA ladder. (B) Agarose gel picture of IEDDA reaction between RNA **13** and Cy5-tetrazine **12**. UV-transilluminator image at (i) 254 nm and (ii) 364 nm. Lane 1: 50 bp DNA ladder.

CONCLUSIONS

We have successfully incorporated the vinyl functionality into RNA by *in vitro* transcription reaction using VUTP **3**. The results demonstrate that VU incorporated into RNA transcripts can serve as a useful handle for chemoselective functionalization of RNA in a modular fashion by oxidative Heck and IEDDA reactions. In particular, the generation of fluorogenic RNA by oxidative Heck reaction with boronic acid/ester is advantageous. It suggests that a screening reaction with appropriate boronic acid/ester substrates could provide direct access to RNA emitting at different wavelengths. Even though conjugation by utilizing oxidative Heck reaction resulted in a mixture of isomers, this study represents a promising initial step on further research on oxidative Heck reaction for the design of made-to-order biologically compatible regioselective and stereoselective ligands for palladium catalysis.^{60,76} Such studies would pave way for selective incorporation of desired biophysical probes, for example, microenvironment sensing probes onto RNA. Furthermore, metabolic labeling of VU followed by posttranscriptional functionalization by these methods could potentially enable the imaging of RNA in cells. Taken together, the studies presented here are expected to complement other bioorthogonal RNA labeling strategies by providing alternative access to RNA labeled with biophysical probes and tags.

ASSOCIATED CONTENT

Supporting Information

The supporting Information is available free of charge in the ACS Publication website.

Experimental procedure, characterization data, Figure S1–S16 and Table S1–S4 (PDF).

AUTHOR INFORMATION

Corresponding Author

*E-mail: srivatsan@iiserpune.ac.in

Notes

The authors declare no competing financial interest.

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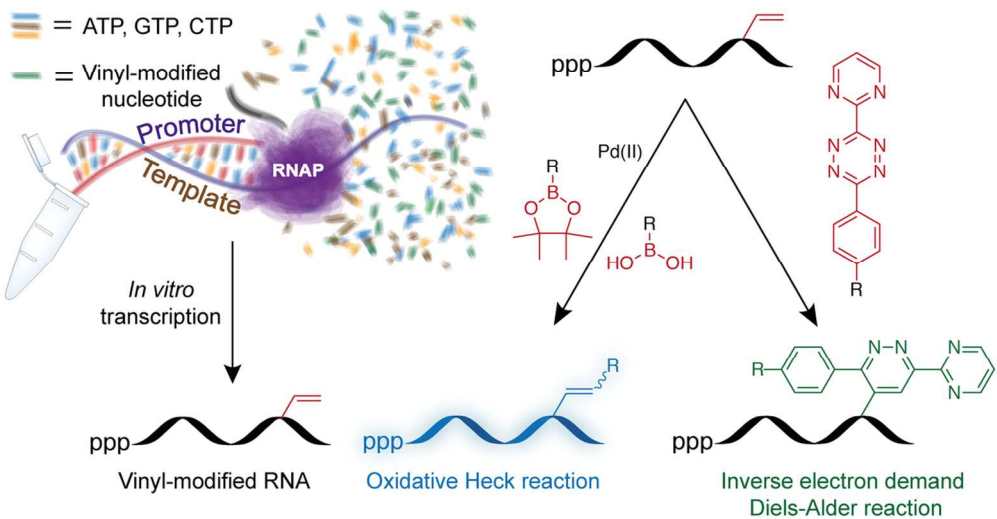
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TOC graphic

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