Biostructural Models for the Binding of Nucleoside Analogs to SARS-CoV-2 RNA-Dependent RNA Polymerase

Andrew J. Prussia* and Spandan Chennamadhavuni

ABSTRACT: SARS-CoV-2 is a positive-sense RNA virus that requires an RNA-dependent RNA polymerase (RdRp) for replication of its viral genome. Nucleoside analogs such as Remdesivir and β-D-N4-hydroxycytidine are antiviral candidates and may function as chain terminators or induce viral mutations, thus impairing RdRp function. Recently disclosed Cryo-EM structures of apo, RNA-bound, and inhibitor-bound SARS-CoV-2 RdRp provided insight into the inhibitor-bound structure by capturing the enzyme with its reaction product: Remdesivir covalently bound to the RNA primer strand. To gain a structural understanding of the binding of this and several other nucleoside analogs in the precatalytic state, molecular models were developed that predict the noncovalent interactions to a complex of SARS-CoV-2 RdRp, RNA, and catalytic metal cations. MM-GBSA evaluation of these interactions is consistent with resistance-conferring mutations and existing structure–activity relationship (SAR) data. Therefore, this approach may yield insights into antiviral mechanisms and guide the development of experimental drugs for COVID-19 treatment.

INTRODUCTION

The pandemic of COVID-19 has brought much of the world to a halt. Its causative agent, SARS-CoV-2, is a betacoronavirus thought to have made an interspecies jump from endemic infection in bats to respiratory infection in humans. From an evolutionary selection viewpoint, the virus is tremendously successful, striking a balance between contagiousness and lethality that has allowed it to escape central China and spread throughout the world. As of today, over 100 million people have been infected and over 2.2 million people have died worldwide. Its human toll has been enormous, and a massive global effort is underway to identify and evaluate vaccines, antiviral agents, and immunotherapies. Although some of the treatment options are now available under emergency use approvals for limited populations, public health efforts are still limited to centuries-old methods of limiting exposure: social distancing, handwashing, protective equipment, and quarantine.

This is not the first coronavirus to threaten mankind. The closest relative to SARS-CoV-2 that causes disease in humans is SARS-CoV, now known as SARS-CoV-1. Its outbreak in 2003–2004 was limited to Asia, but the fatality rate was much higher, around 10%. There have been several attempts to develop an antiviral drug for SARS-CoV-1, and while some showed promise, without an ongoing outbreak, no treatment was finalized. Unlike the positive-sense RNA virus SARS-CoV-2, Ebolavirus is a negative-sense RNA virus but went through a similar antiviral development cycle: without the pressure from an uncontrollable spread, no antiviral drugs were approved. Fortunately, these efforts did leave us with late-stage experimental drugs that could be repurposed against SARS-CoV-2. One class of these repurposed antiviral drugs that have shown activity against SARS-CoV-2 are nucleoside analogs (Figure 1). Coronaviruses, Filoviruses (such as Ebola virus) and other single-stranded RNA viruses require an RNA-dependent RNA polymerase (RdRp) for replication of

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their genomes. The catalytic cores of these enzymes are remarkably conserved across the virus families since the basic functions of holding viral RNA, binding an incoming ribonucleotide, and catalyzing the polymerase reaction are necessary steps in all these viral families. It may even be possible for a nucleoside analog to have activity against multiple viral families and thus function as frontline therapy in the next outbreak.\textsuperscript{13,14}

Although the RdRp of SARS-CoV-2 is an attractive target, effective inhibition by nucleoside analogs requires a complicated sequence of metabolic transformations because the active agent is not the nucleoside analog. It is instead the nucleoside triphosphate (NTP) that is needed for its metabolic activation.

Figure 1. (a) Nucleoside analogs with known antiviral activity against SARS-CoV-2 are shown in their dosed and metabolically active forms. (b) Commercially available compounds 1, 2, and 3 are predicted to be RdRp inhibitors in their triphosphorylated form.
incorporation into the viral RNA. Such a charged species, however, would have limited absorption and distribution across cellular membranes.\(^7\text{-}^\text{11}\) As a consequence, nucleoside drugs are either dosed as nucleoside analogs (i.e., the compound possesses a free 5′-hydroxyl) or as monophosphate prodrugs, both of which typically possess good cellular permeability. In the former case, three different cellular kinases sequentially phosphorylate the 5′-hydroxyl to produce the desired NTP. In the latter case, endogenous enzymes (esterases, phospholipases, phosphoramidases, etc.) unravel the prodrug to reveal the nucleoside monophosphate. The first phosphorylation step, which is often rate-limiting, is then followed by kinetically favored second and third phosphorylation steps. In order for the NTP to inhibit viral replication, it must inhibit the RdRp from functioning. If the NTP–RdRp interaction is favorable, it will be incorporated into a growing strand of RNA and either terminate its elongation or cause it to become dysfunctional. All these steps are necessary but not sufficient for the nucleoside analog in question to be a viable antiviral agent. The final hurdle relates to the safety of the nucleoside analog, i.e., it must exhibit a binding preference at the viral RdRp over cellular nucleotidyl receptors.

As there was no structural information available, we initially built a homology model for SARS-CoV-2 RdRp based on structural information for other positive-sense, single-strand RNA viruses. Recently, however, cryo-electron microscopy structures have been reported for coronaviruses: an apo RdRp structure (PDB code 6M71),\(^15\) an RdRp with duplex RNA bound (PDB code 6YYT),\(^16\) and an RdRp structure covalently bound to Remdesivir (PDB code 7BV2),\(^17\) which dramatically advanced our knowledge of the SARS-CoV-2 RdRp complexes. The three studies found structures that closely matched the SARS-CoV-1 RdRp structure (PDB code 6NUR) published last year, which is not surprising given their 96% sequence identity.\(^18\) However, apo RdRp structures are of limited utility for the binding rationalization of nucleoside analogs that require base-pairing to the template RNA strand to function as inhibitors. Likewise, the RNA-bound and Remdesivir-bound structures are also not ideal because both are reaction products of enzymatic catalysis and not the precatalytic state where nucleoside analogs must compete with endogenous nucleotides as substrates for the polymerization reaction. Multiple structural features differ between the Remdesivir structure and initialization complexes for other viral RdRps including the positions of catalytic metals, aspartate side chains, and the formation of a loop interacting with template RNA (Figure 2a–c).\(^19\text{-}^\text{23}\)

Ahmad and colleagues used homology modeling and other computational techniques with the structures described above to investigate possible small-molecule inhibition of the SARS-CoV-2 RdRp initiation complex with GTP.\(^22\) Although this approach may provide inhibitors for the initiation stage without primer-bound RNA, the primary inhibitory mechanism by Remdesivir and other nucleoside analogs is delayed chain termination at the \(i + 3\) position.\(^23\text{-}^\text{24}\) Gordon et al.\(^25\) demonstrated this mechanism through enzyme kinetics and developed a structural model for noncovalent interaction of Remdesivir but did not rationalize existing SAR data. Independently, we built precatalytic biostructural models for SARS-CoV-2 RdRp to evaluate the binding of Remdesivir and many other antiviral drugs of SARS-CoV-2 RdRp such as β-D-N′-hydroxycytidine,\(^13\text{-}^\text{25}\) Favipiravir,\(^26\) and other potential nucleoside analog inhibitors.\(^27\) These models were developed with the complementary template RNA nucleobases to accommodate nucleoside analogs with different nucleobases. A mutated version of SARS-CoV-2 RdRp with resistance to Remdesivir was also developed. Using these models, the potential binding of phosphorylated nucleoside analogs was evaluated using the MM-GBSA protocol and compared with the binding of natural RNA nucleotides. The obtained results show that the models are beneficial in developing the next generation of nucleoside analogs against SARS-CoV-2.

![Figure 2](https://doi.org/10.1021/acscim.0b01277)

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Development of SARS-CoV-2 RdRp-RNA-Metal Complexes. The 2.5 Å resolution electron microscopy structure of the SARS-CoV-2 nsp12-nsp7-nsp8 (PDB code 7BV2, Figure 2a) was aligned with the 1.8 Å X-ray crystallography structure of the Norwalk virus polymerase (PDB code 3BSO, Figure 2b) using the Protein Structure Alignment tool by Schrödinger, LLC. This tool aligns the secondary structure features that are common to the two proteins. The overall RMSD in C-alpha positions was 3.2 Å with an alignment score of 0.47 (alignment scores <0.6 indicate good alignment; proteins with >0.8 scores may not be similar structurally). Although the sequence homology between these two proteins is low with this alignment (8% identity, 19% similarity), the binding site residues are conserved with residues within 5 Å having an 1.6 Å RMSD in their C-alpha atom positions and the conserved viral RdRp motifs A, B, C, and D overlapping (the E and F motifs are not available in the 3BSO structure). To create the precatalytic model, Remdesivir and the catalytic metal cations of 7BV2 were replaced with ATP and the catalytic metal cations of the aligned structure of 3BSO. The resulting structure was prepared for molecular calculations and evaluated for steric clashes and other issues by the Protein Preparation Wizard. With the geometry of ATP, RNA, and metal cations fixed in a precatalytic state, loop refinement of residues forming the 676-686 loop and side-chain predictions of all residues were performed with Prime using the VSGB continuum solvation model for water and the OPLS3 force field. Extended serial loop sampling was chosen during loop refinement to ensure that sufficient sampling was performed to identify the lowest-energy loop conformation. We hypothesized that optimizing the enzyme around a precatalytic state of its substrate would shift its conformation to a state suitable for nucleoside analog binding.

Optimization of Nucleoside Analogs with the MM-GBSA Protocol. With the structure optimized toward a precatalytic state with ATP bound, other nucleoside analogs were modeled by modifying the ATP ribose and base rings to the desired substituents. To model GTP, CTP, UTP, and the nucleoside analogs that mimic these bases, the template RNA base was mutated to the complementary base. The modified structures were created in chemically reasonable conformations to avoid high-energy contacts upon optimization. Optimization calculations were performed with the Prime MM-GBSA protocol using the same solvation and force field settings as above. The ligand (ATP, GTP, CTP, UTP, or the nucleotide analogs) was allowed complete flexibility, while the protein, RNA, and metal cations within 6 Å of the ligand were flexible with harmonic constraints of 1.0 kcal/mol-Å². The difference in MM-GBSA ΔG (NS) binding energies (∆∆Gbind) was calculated by subtracting the binding energy of the natural nucleotide (that expected to be displaced in the active site) from the binding energy of the nucleoside analog of interest.

Density Functional Theory Calculations for Tautomers of EIDD-1931. The energy difference between the enamine and oxime forms of EIDD-1931 was calculated on N-methyl model systems of the nucleobase by optimization using Jaguar at the M06-2X/cc-pVTZ(-f) level with the Poisson–Boltzmann continuum water solvation model. The N-methyl model system of the nucleobase was used to (1) reduce calculation time, (2) remove alternative conformations, which would confound the final energy but not the tautomization equilibrium, and (3) focus on subtle differences in the energetics of tautomers in the solution phase. The stereoelectronic effect of the sugar or the triphosphate group is expected to be negligible in the tautomization equilibrium of the binding conformation of EIDD-1931.

RESULTS AND DISCUSSION

The starting point for building this model was the 2.8 Å resolution electron microscopy structure of the SARS-CoV-2 nsp12-nsp7-nsp8 with Remdesivir covalently bound (Figure 2a). This structure was modified into a precatalytic state by removing Remdesivir from the primer RNA and inserting adenosine triphosphate (ATP) into the state captured by the 1.7 Å resolution X-ray crystallography structure of the Norwalk virus RdRp initiation complex (PDB code 3BSO, Figure 2b). The Norwalk virus RdRp structure was chosen because of its structural similarity to SARS-CoV-2 and its unique capture of the triphosphate form of a nucleotide instead of the diphosphate form of HCV (Figure 2c). The two catalytic metals were also shifted to the position found in the Norwalk virus RdRp. Once in place, a slight shift in the 676-686 loop was necessary to accommodate the nucleotide in this state. This was remedied by a loop prediction algorithm and resulted in a subtle conformational change of this loop. The protein side chains were also optimized to this form by a side-chain prediction algorithm, which converged on a low-energy conformation that mimics the precatalytic state found in the Norwalk and Hepatitis C Virus RdRp structures (Figure 2c). Similar side-chain rotamers and loop conformations indicate that these refinement procedures have shifted the enzyme from a post- to a precatalytic form observed for the Norwalk virus RdRp. An overview of the modeling process is illustrated in Figure 2d. Our biostructural model would be extremely beneficial in prioritizing novel nucleoside analogs, while structural biologists experiment with methodologies to capture the SARS-CoV-2 RdRp in its precatalytic state using X-ray crystallographic or cryo-electron microscopy methods.

The energetics of the resulting complex can be interrogated by the MM-GBSA binding energy procedure. Although other computational methods such as free energy of perturbation (FEP) or hybrid QM/MM could be applied, our focus was on (a) maintaining the proper geometry for the polymerase reaction and (b) a method that could be applied rapidly to many nucleoside analogs. This is a reasonable approach to predict inhibition of nucleoside analogs, considering other inherent variables such as phosphorylation and ADMET. This protocol allowed the nucleotide to freely optimize its structure in the presence of the protein, duplex RNA, and metal cations. If there were unfavorable interactions between this substrate and the surrounding atoms, it was anticipated that the structure might become distorted or the nucleotide would be pushed out of the catalytic binding pocket. However, with this approach, the nucleotide maintained its conformation, shifting slightly in position but maintaining base-pair interactions with the template strand of RNA, indicating that its binding pose may be similar to the initial stage of the polymerase reaction (Figure 2c). The metal cations also shifted slightly in their position while maintaining coordination interactions with the catalytic aspartate side chains. The MM-GBSA protocol provided a calculated binding interaction energy. Because this energy is dominated by the electrostatic term from the charged interactions between the
Table 1. Nucleoside Anti-Coronavirus Activity and Predicted Binding in the Active Site of the SARS-CoV-2 RdRp Model Complex

| Nucleoside analog                  | Binding pose in SARS-CoV-2 RdRp model complex | Antiviral activity          | $\Delta G_{\text{bind}}$ (kcal/mol) |
|------------------------------------|------------------------------------------------|-----------------------------|-------------------------------------|
| Remdesivir triphosphate            | [Image]                                         | 0.77 μM: SARS-CoV-2 EC50  
0.069 μM: SARS-CoV-1 IC50  
0.074 μM: MERS-CoV IC50         | -13.6                             |
| EIDD-1931 triphosphate             | [Image]                                         | 0.08 μM: SARS-CoV-2 IC50  
0.14 μM: SARS-CoV-1 IC50  
0.024 μM: MERS-CoV IC50         | -21.8                             |
| EIDD-1931 triphosphate,            | [Image]                                         | 57.7 μM: SARS-CoV-1 EC50  
68.4 μM: MERS-CoV-1 EC50         | -5.5                              |
| (oxime tautomer)                   |                                                 |                            |                                     |
| Favipiravir triphosphate           | [Image]                                         | ATP mimic 118.3 μM: SARS-CoV-2 IC50 [CPE]  
207.1 μM: SARS-CoV-2 IC50 [replication]          | -12.7                             |
| GTP mimic                          |                                                 |                            | -1.0                               |
| Galidesivir triphosphate           | [Image]                                         | 0.03 μM: HCoV-NL63 IC50        | -5.73                              |
| 6-Azauridine triphosphate          | [Image]                                         | 0.03 μM: HCoV-NL63 IC50        | -5.73                              |
| Acyclovir Fleximer triphosphate    | [Image]                                         | 23 μM: MERS-CoV IC50  
8.8 μM: HCoV-NL63 IC50       | +2.5                               |
| Ribavirin triphosphate             | [Image]                                         | 50-819 μM: SARS-CoV-1 IC50  
20 μM: MERS-CoV IC50           | +6.0                               |
| Gemcitabine triphosphate           | [Image]                                         | 1.22 μM: MERS-CoV IC50  
4.96 μM: SARS-CoV-1 IC50       | -1.56                              |
| Mleziribine triphosphate           | [Image]                                         | 13.5 μM: MERS-CoV IC50  
62 μM: SARS-CoV-1 IC50        | +13.5                              |
| Commercial compound 1              | [Image]                                         | Not yet available             | -13.5                              |
| Commercial compound 2              | [Image]                                         | Not yet available             | -16.8                              |
| Commercial compound 3              | [Image]                                         | Not yet available             | -16.3                              |

*Energies described by $\Delta G_{\text{bind}}$ are relative to the natural nucleotide predicted to be displaced by the nucleoside analog. Antiviral efficacies against SARS-CoV-2, SARS-CoV-1, MERS-CoV, and HCoV-NL63.*

nucleotide phosphates, catalytic metals, and negatively charged aspartate side chains (the Coulombic term is >3 times the vDW term but countered by the solvation term), it is not expected to give an accurate absolute binding energy that would be
correlated with antiviral activity. It could, however, provide relative energies to compare the binding of nucleotide analogs with the same nucleobase or bioisostere. This is described as \( \Delta G_{\text{bind}} \), which compared the binding energy of a nucleotide analog with its respective natural nucleotide (Table 1). Because this competitive binding is one of the many conditions necessary to achieve viral inhibition, this energetic comparison is unlikely to be correlated with antiviral activity but could provide a basis for SAR comparison between analogs that could classify analogs into three categories: (a) likely RdRp inhibitors, (b) potential RdRp inhibitors, and (c) unlikely RdRp inhibitors. This virtual evaluation approach might be advantageous in prioritizing which nucleosides to screen for their antiviral efficacy, given the limited synthesis and virology resources.

Remdesivir (Veklury) is the first FDA-approved drug for Covid-19 treatment.\(^\text{35}\) It was discovered and developed by Gilead Pharmaceuticals and has completed eight phase 3 clinical trials. Currently in 49 active or recruiting phase 3 clinical trials, it is a phosphoramidate prodrug C-nucleoside analog with both a modified nucleobase and ribose sugar (Figure 1).\(^\text{37,38}\) Its base mimics the hydrogen bond pairing of adenosine, but its ribose has a 1′-nitrile substituent distinct from the natural nucleotide. To dock its respective nucleotide into the SARS-CoV-2 RdRp complex, ATP was modified in place to Remdesivir and the complex was subjected to MM-GBSA minimization. The resulting structure showed a slight shift in position but established a polar contact between the 1′-nitrile substituent and Ser759 while maintaining strong base-pairing hydrogen bonds with the template RNA nucleobase (Table 1). Notably, the 1′-nitrile interaction was different in the covalently bound Remdesivir structure, which had a similar base pairing to Thr687. Compared with ATP, Remdesivir had a more favorable MM-GBSA binding energy by \(-13.1\) kcal/mol. This is consistent with Remdesivir's high level of antiviral activity and kinetic assays, which show that the enzyme is more selective for Remdesivir over ATP.\(^\text{23}\)

The ribonucleoside analog β-D-N3′-hydroxycytidine (NHC, EIDD-1931) is only modified from cytidine by an oxime at the 4 position of the nucleobase (Figure 1),\(^\text{25}\) yet it showed antiviral efficacy against a panel of viruses, including SARS-CoV-2. Its S′-isopropyl ester prodrug, Molnupiravir (EIDD-2801) was found to have better ADMET and is orally bioavailable.\(^\text{13}\) Currently, the drug is in two phase 3 clinical trials (NCT04575584 and NCT04575597). The oxime also tautomerizes to the enamine form. Interestingly, this provides the analog with two base-pairing opportunities: the oxime tautomer mimics uridine, while the enamine imitates cytidine.\(^\text{39}\) Optimization of both tautomer nucleobases at the M06-2X/cc-pVTZ(-f) level with the Poisson–Boltzmann continuum water solvation method indicated near equivalent populations of both tautomers. It was found that the oxime form is slightly favored by 0.2 kcal/mol in the solution phase (although the tautomization equilibrium could be shifted upon binding with the enzyme). Analyses of the active triphosphorylated forms of both tautomers in the SARS-CoV-2 RdRp model complex revealed nearly identical positioning to natural uridine and cytidine, but with an additional hydrogen-bonding interaction between the enamine or oxime hydroxyls with the preceding base in the primer RNA strand (Table 1). The MM-GBSA protocol calculated binding energies \(-21.8\) and \(-10.1\) kcal/mol relative to the binding of unmodified uridine and cytidine, respectively, consistent with the potent broad-spectrum antiviral efficacy of this nucleoside analog. This also indicates that nucleoside analogs to bases other than ATP can be evaluated in this model system. It is important to note that although both the DFT and the MM-GBSA calculations give energy results in kcal/mol, the MM-GBSA calculations are based on molecular mechanics and are not expected to give thermodynamically accurate energies in this system.

Favipiravir (Figure 1) is unique from all other nucleoside analogs discussed earlier as it is dosed in substituted pyrimidine form. Once in the cell, Favipiravir undergoes phosphoribosylation to the active antiviral form Favipiravir ribofuranosyl-5′-triphosphate.\(^\text{27}\) The modified pyrimidine ring of Favipiravir is a unique bioisostere of adenosine or guanosine in its base-pairing hydrogen bonding capability. In the SARS-CoV-2 RdRp model complex, Favipiravir RTP retained a similar position to either natural adenosine or guanosine nucleotides but formed more favorable interactions when acting as an adenosine mimic (Table 1). The binding energy relative to adenosine is \(-12.7\) kcal/mol, suggesting that binding to the SARS-CoV-2 RdRp could be favorable. Favipiravir does in fact inhibit SARS-CoV-2 through this mechanism with an IC\(_{50}\) of 118.3 \(\mu\)M, as was recently reported.\(^\text{28}\) This level of inhibition may be due to phosphoribosylation steps, which could limit the supply of the active ribofuranosyl-5′-triphosphate form.

Galidesivir (BCX4430) (Figure 1) is currently being evaluated in the clinical trials for COVID-19 (NCT03891420) by BioCryst Pharmaceuticals. This experimental drug is an adenosine nucleoside analog, containing a slightly modified base and an azasugar. It has broad-spectrum activity against nine virus families with 57.7 and 68.4 \(\mu\)M EC\(_{50}\) against SARS-CoV-1 and MERS-CoV, respectively.\(^\text{40}\) When docked in the SARS-CoV-2 RdRp complex, it is predicted to have moderately favorable interaction based on a \(\Delta G_{\text{bind}}\) energy of \(-5.5\) kcal/mol. Thus, relative binding energies in the range of \(-5\) to \(-10\) kcal/mol may represent analogs with the modest activity.

6-Azauridine (Figure 1) has 0.03 \(\mu\)M EC\(_{50}\) activity against a human coronavirus NL63 (HCoV-NL63), another pathogenic virus associated with acute respiratory illness.\(^\text{41}\) The overall sequence similarity between the RdRps of HCoV-NL63 and SARS-CoV-2 is moderate (37% identity, 53% homology), so a common inhibition is possible. Evaluation in the model of SARS-CoV-2 RdRp predicts an interaction similar to Galidesivir, with a \(\Delta G_{\text{bind}}\) energy of \(-5.7\) kcal/mol.

The model did not predict favorable binding for an Acyclovir Fleximer, while this nucleoside analog has 23 and 8.8 \(\mu\)M EC\(_{50}\) activities against MERS-CoV and HCoV-NL63, respectively.\(^\text{41}\) This biaryl compound was designed to not only mimic a pyrimidine base pairing pattern but also be able to adopt alternative binding modes.\(^\text{42}\) In the model, the compound was only able to fit in the catalytic site as an adenosine mimic, base pairing with the uracil base of the template RNA, but it is possible that the compound may function as another nucleobase mimic or adopt an alternative binding mode. This compound has not yet been shown experimentally to function as a chain terminator (either immediate or delayed) of RNA replication, so it may function by a mechanism that does not require RNA incorporation. Since our models are based on competition with natural nucleotides in a precatalytic state, it might not be able to predict other inhibition mechanisms. An additional kinetic
assay or structural information for this compound may be necessary to elucidate its inhibition mechanism.

Ribavirin is another nucleoside analog with broad-spectrum activity but is thought to act pleiotropically by inhibiting the host inosine monophosphate dehydrogenase, increasing expression of interferon-gamma and tumor necrosis factor-alpha and increasing the frequency of viral mutations. Only the latter mechanism would require direct binding of Ribavirin triphosphate with the SARS-CoV-2 RdRp complex, but the effect could be negated by the exoribonuclease proofreading function of beta coronaviruses. In SARS-CoV-1, Ribavirin triphosphate is known to be incorporated, but then excised by this function, making this mechanism unlikely to play a role in ribavirin inhibition of the SARS-CoV-2 virus. In an enzyme kinetics study, the $K_m$ for Ribavirin triphosphate binding is 136 $\mu$M for SARS-CoV-1 RdRp, 4-fold higher than Remdesivir triphosphate. The $\Delta\Delta G_{bind}$ energy for Ribavirin triphosphate is +6.0 kcal/mol, indicating that the model may not identify weak binding interactions even if the nucleotide analog is capable of RNA incorporation.

Mizoribine and Gemcitabine also are active against coronaviruses, SARS-CoV and MERS-CoV, at micromolar levels. Gemcitabine is only predicted to have a slightly favorable interaction in the SARS-CoV-2 model, with a $\Delta G_{bind}$ energy of $-1.56$ kcal/mol, while Mizoribine is disfavored by $13.5$ kcal/mol. The mechanism of viral inhibition is not confirmed for either of these nucleoside analogs: Gemcitabine is a chemotherapy drug and Mizoribine is an immunosuppressant, so it is possible that their antiviral activities are due to host-directed inhibition rather than direct interaction with a viral protein. Figure 3 summarizes the relationship between coronavirus inhibition from SARS-CoV-2, SARS-CoV-1, or MERS-CoV-1 and binding energies in the model (the coronavirus most similar to SARS-CoV-2 with available inhibition data is shown). Nucleoside analogs with documented RdRp inhibition are predicted to have favorable binding relative to the natural nucleotide. Analogs with $\Delta G_{bind}$ energies more favorable than $-10$ kcal/mol are often active at sub-micromolar levels against coronaviruses. The model may therefore be useful to prioritize synthetic and biological resources for developing nucleoside analogs against coronaviruses, along with the necessary considerations of ADMET properties, phosphorylation, and synthetic feasibility. The complex cellular and viral interactions necessary for nucleoside analog inhibitions preclude a quantitative prediction of antiviral activity using this model, but the model may serve to classify nucleoside analogs as probable, possible, or unlikely SARS-CoV-2 RdRp inhibitors. To demonstrate the utility of this approach, commercially available nucleoside analogs from the Granlen library (Granlen, Inc.) were evaluated in the models. Examples of likely RdRp inhibitors are shown in Table 1. The model predicts that 2’-methoxy substitution (Commercial Compound 1) and iodo-substitution at the 5-position of the cytosine nucleobase (Commercial Compound 2) are favorable ($\Delta G_{bind}$ energies of $-13.5$ and $-16.8$ kcal/mol, respectively). Although both substitutions appear to be compatible in one compound (Commercial Compound 3), there seems to be no apparent additive effect on the binding energy ($\Delta G_{bind}$ energy of $-16.3$ kcal/mol). These and other commercially available nucleoside analogs, as well novel entities, continue to be investigated by our research group.

Mutations conferring resistance are a hallmark of RNA viruses under antiviral pressure, even with coronaviruses that also have an exonuclease proofreading functionality. The two mutations that have been observed against Remdesivir in SARS-CoV-1 are Phe480Leu and Val557Leu and result in a 5.6-fold decrease in inhibition. These mutations were modeled in the SARS-CoV-2 RdRp model and the complex subjected to the side-chain prediction of all protein residues. The complex converged on an almost identical arrangement of side chains (Figure 4). Subjecting Remdesivir to the MM-GBSA protocol in the mutated model did not cause a significant difference in the binding pose but did reduce $\Delta G_{bind}$ to $-0.3$ kcal/mol. It appears that extending the isopropyl group of Val557 to isoleucine forces this nonpolar residue near the more polar aromatic ring of Remdesivir. The Phe480Leu mutation is distal to the active site and RNA binding residues. This mutation may play a role in the dynamics of the polymerase as it opens and closes in the process of binding RNA and catalyzing the polymerase reaction. Resistance-conferring mutations to Remdesivir have not yet been observed in SARS-CoV-2 RdRp but are likely to occur through the same mechanisms based on the 96% sequence identity between the RdRps of these viruses. Interestingly, EIDD-1931 maintains its activity against a coronavirus RdRp bearing these mutations. Its $\Delta G_{bind}$ in the enzyme tautomer reduced to $-5.6$ kcal/mol but was still in the range of possibly active nucleoside inhibitors similar to Galidesivir.

CONCLUSIONS

The development of biostuctural models for SARS-CoV-2 RdRp has enabled the virtual evaluation of nucleoside analogs for antiviral activity. Known nucleoside analog inhibitors, such as Remdesivir and Molnupiravir, are predicted to bind favorably in the precatalytic models of the RdRp, demonstrating that multiple nucleobase and ribose substitutions can be evaluated by the models with correct activity classification. Models with mutations that confer resistance to Remdesivir are able to quantify a structural rationale for this effect that is consistent with the absence of resistance to Molnupiravir. With these tools, commercially available nucleoside analogs with potential anti-coronavirus activity were identified and

Figure 3. Graph of coronavirus inhibition of nucleoside analogs versus their $\Delta G_{bind}$ energies evaluated in the SARS-CoV-2 RdRp model. Blue, red, and black circles indicate SARS-CoV-2, SARS-CoV-1, and MERS-CoV-1 inhibition activities, respectively. Coronavirus activities on the y-axis are log-scaled.

| Compound       | Likely RdRp inhibitor | Potential RdRp inhibitor | Unlikely RdRp inhibitor |
|----------------|-----------------------|--------------------------|------------------------|
| Favipiravir(A) | ✓                     | ✗                        | ✗                      |
| Remdesivir     | ✓                     | ✓                        | ✗                      |
| EIDD-1931(U)   | ✓                     | ✓                        | ✓                      |
| 6-AzaTudine    | ✓                     | ✓                        | ✓                      |

Table 1. Examples of likely, potential, and unlikely SARS-CoV-2 RdRp inhibitors.
prioritized for testing. The authors will continue these efforts to aid other researchers in the development of nucleoside analogs against the global pandemic of COVID-19. All modeled structures described here are available for download in Supporting Information 1–4. While safe and effective (>95%) vaccines (Pfizer, Moderna, etc.) against SARS-CoV-2 are now available, inoculating the world population is a costly, time-taking endeavor. Limited vaccine availability, logistics in distribution, compliance issues, and emerging variants of viruses due to mutations mean that novel direct-acting antivirals will still be necessary to tackle this ongoing pandemic as well as threats that mankind might face in the future. The authors aspire that these models, along with efforts to develop different classes of antivirals, vaccines, and other therapies, will facilitate a global solution to the current crisis.

ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.jcim.0c01277.
- PDB files for the optimized nucleoside analog complexes for atp (ZIP)
- PDB files for the optimized nucleoside analog complexes for ctp (ZIP)
- PDB files for the optimized nucleoside analog complexes for utp (ZIP)
- PDB files for the optimized nucleoside analog complexes for gtp (ZIP)

AUTHOR INFORMATION

Corresponding Author
Andrew J. Prussia — Department of Chemistry, Emory University, Atlanta, Georgia 30322, United States; orcid.org/0000-0002-7038-9520; Email: aprussia@me.com.

Author
Spandan Chennamadhavuni — Department of Chemistry, Emory University, Atlanta, Georgia 30322, United States; orcid.org/0000-0002-1026-9527

Complete contact information is available at: https://pubs.acs.org/10.1021/acs.jcim.0c01277

Notes
The authors declare no competing financial interest.

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