

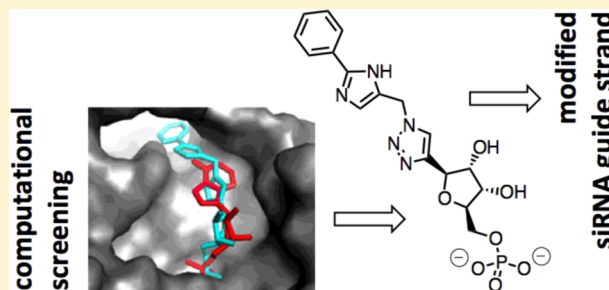
Short Interfering RNA Guide Strand Modifiers from Computational Screening

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S Supporting Information

ABSTRACT: Short interfering RNAs (siRNAs) are promising drug candidates for a wide range of targets including those previously considered “undruggable”. However, properties associated with the native RNA structure limit drug development, and chemical modifications are necessary. Here we describe the structure-guided discovery of functional modifications for the guide strand 5′-end using computational screening with the high-resolution structure of human Ago2, the key nuclease on the RNA interference pathway. Our results indicate the guide strand 5′-end nucleotide need not engage in Watson–Crick (W/C) H-bonding but must fit the general shape of the 5′-end binding site in MID/PIWI domains of hAgo2 for efficient knockdown. 1,2,3-Triazol-4-yl bases formed from the CuAAC reaction of azides and 1-ethynylribose, which is readily incorporated into RNA via the phosphoramidite, perform well at the guide strand 5′-end. In contrast, purine derivatives with modified Hoogsteen faces or N2 substituents are poor choices for 5′-end modifications. Finally, we identified a 1,2,3-triazol-4-yl base incapable of W/C H-bonding that performs well at guide strand position 12, where base pairing to target was expected to be important. This work expands the repertoire of functional nucleotide analogues for siRNAs.



INTRODUCTION

Short interfering RNA (siRNA)-triggered gene knockdown via the RNA interference (RNAi) pathway has revolutionized the study of gene function and spurred the development of new oligonucleotide-based therapeutics.^{1,2} However, the natural RNA structure poses limits to drug development, and modifications are necessary to improve properties (e.g., increase potency, reduce nuclease sensitivity, facilitate delivery, reduce off-target effects, etc.).^{3–6} Thus, it is important to identify modifications at specific positions in the siRNA strands where changes in the natural RNA structure improve siRNA performance.^{7,8} The key step in the RNAi pathway is cleavage of target RNA by the RNaseH-like enzyme Ago2 in complex with the siRNA guide strand. Recent mechanistic⁹ and structural studies^{10,11} have shown that the four domains of the human Ago2 protein (N-terminal, PAZ, MID, and PIWI) each have a distinct role in guide strand binding and the target RNA cleavage reaction (Figure 1). For instance, the hAgo2 PAZ domain binds the 3′ end of the guide strand with direct contacts made to the last three nucleotides.^{11,12} On the other hand, nucleotides 2–6 of the guide strand are bound to the PIWI domain in an A-form helical geometry with their Watson–Crick (W/C) edges available for binding target RNA. The PIWI domain is also responsible for the target RNA cleavage reaction. Importantly, the 5′-most nucleotide (i.e., guide nucleotide 1) is tucked into a MID domain pocket where extensive contacts are made to both the 5′-phosphate and the nucleobase (Figure 1). Interactions with the edge of the

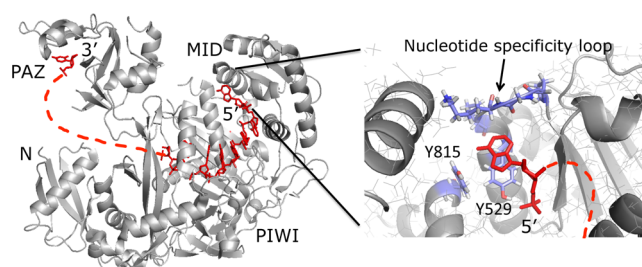


Figure 1. hAgo2 interactions with an RNA guide strand.^{10,13}

base via a rigid loop in the protein (i.e., the nucleotide specificity loop) explain the MID domain's preference for binding pU and pA over pG and pC (Figure 1).¹³ This nucleotide does not W/C pair with target RNA, but modifications to its structure that reduce MID domain binding affinity reduce siRNA efficacy.¹⁴ For instance, an abasic site at this location results in weaker MID domain binding and lower target cleavage activity.⁹ This is most likely because the nucleobase at guide position 1 stacks onto Tyr 529 of the MID domain and an abasic residue lacks this interaction (Figure 1). Since this guide strand nucleotide is not involved in target nucleotide recognition yet influences siRNA activity, modifications at this position have the potential to enhance RNAi

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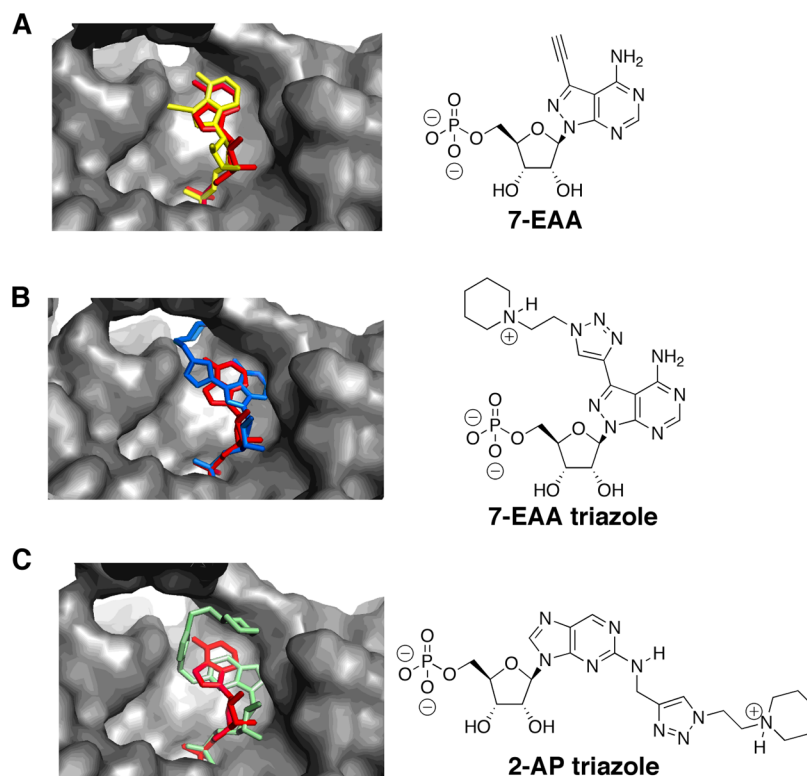


Figure 2. Predicted hAgo2 binding modes for purine analogues positioned at a guide strand 5'-end. Each analogue is shown in overlay with adenosine (red) in the guide strand 5'-end binding site of hAgo2 shown as a gray surface.¹⁰ (A) 7-EAA (yellow), (B) 7-EAA triazole (blue), and (C) 2-AP triazole (green).

potency and could be generally useful for all siRNAs. Furthermore, two recent reports of crystal structures of human Ago2-guide strand complexes clearly define 5'-end/hAgo2 interactions and suggest a structure-guided approach may be possible for the discovery of functional 5'-end modifications.^{10,11} We report here a combined computational screening and synthesis strategy to identify new 5'-end modifications for siRNA guide strands. We used the structure of a hAgo2/guide strand complex and computational screening of nucleobase analogues accessible by copper-catalyzed azide/alkyne cycloaddition (CuAAC) reactions. The screening identified several high scoring candidates for 5'-end modifiers including triazole derivatives that appeared capable of accessing a cleft in hAgo2 adjacent to the 5' nucleotide-binding site. Synthesis and testing of a group of guide strand 5'-end modifications showed a good correlation between siRNA potency and predicted hAgo2 interactions.

RESULTS AND DISCUSSION

Computational Screening for 5'-End Modifiers. To access the computationally predicted hAgo2 binding affinity of guide strand 5'-terminal nucleobases, we constructed a library of synthetically accessible structures (see below), expanded this library to conformers of each molecule, and then docked the conformational library into a rigid receptor created from the published crystal structure of hAgo2 bound to guide strand RNA (4EI1).¹⁰ This was achieved using the *OpenEye* Suite of programs (see Experimental Section).¹⁵ The receptor was created in such a way as to ensure that even the largest analogues investigated could be accommodated and with constraints imposed on the 5'-phosphate group and the 3'-hydroxyl to simulate how the 5'-end nucleotide of the guide

strand is bound by hAgo2. Each base was assigned a binding score measuring the quality of the fit of the analogue into the binding pocket (e.g., π stacking, sterics, complementarity of H-bonding donors and acceptors, etc.). These scores were generated using several different computational docking scoring methods as well as analysis of the top pose overlay with the guide strand 5'-end adenosine from the hAgo2-RNA crystal structure. A full description of the docking and scoring method can be found in the Experimental Section.

Our lab has prepared adenosine analogues that direct substituents into either the minor^{8,16,17} or major groove¹⁸ when present in duplex RNA. Since adenine (along with uracil) is a preferred natural nucleobase ligand for the Ago2MID domain, we initially screened a library of adenosine analogues for good hAgo2-binding scores as described above.¹³ For instance, 7-ethynyl-8-aza-7-deazaadenosine (7-EAA) is readily incorporated into RNA via the phosphoramidite and serves as a precursor to 7-triazole derivatives by CuAAC reactions (Figure 2).¹⁸ However, 7-EAA was predicted to fit poorly into the 5'-nucleotide binding pocket of hAgo2 (Figure 2) and did not score well relative to adenosine in the computational screening (Table 1). Indeed, the lowest-energy binding pose for 7-EAA has the ethyne clashing with the aromatic ring of Tyr 815 of the adjacent PIWI domain (Figure 1, Figure 2A). Triazoles formed from reaction of this analogue with different azides were also screened (57 total) (see Supplementary Figure 1 and Supplementary Table 1) and generally had poor scores and inferior fits in the binding site compared to adenosine (see 7-EAA triazole in Figure 2B, Table 1).

2-Aminopurine analogues can be used in duplex RNA as adenosine replacements that project substituents into the minor groove.^{8,16} Our earlier work showed that an *N*-ethylpiperidine

Table 1

	predicted hAgo2 binding ^a	siRNA activity ^b
adenosine	1.0	++
7-EAA	8.8	+
7-EAA triazole	9.2	+
2-AP-triazole	9.5	+

^aScores normalized from 1 to 10 from several scoring methods (see Experimental Section). A lower number represents better-predicted binding to guide strand 5'-end. ^bExperimental siRNA activity determined at 100 pM. +++ = <10% luciferase activity remaining after knockdown; ++ = 10–40%; + = 41–70%; – = >70% (Figure 3).

triazole derived from 2-propargylaminopurine (i.e., 2-AP-triazole) is well-tolerated at position 14 of the guide strand but not at position 2 (Figure 2C).⁸ When evaluated computationally for binding into the hAgo2 guide strand 5'-end binding site, this structure also did not fit well and scored poorly relative to adenosine (Figure 2C) (Table 1).

These computational screening results suggested that the adenosine analogues would likely inhibit RNAi activity when positioned at the guide strand 5'-end. To determine the effect of the 2-AP triazole at this position, it was incorporated into an siRNA that targets the PIK3CB sequence present on a luciferase reporter plasmid and activity for the modified siRNA was measured in HeLa cells (Figure 3). Consistent

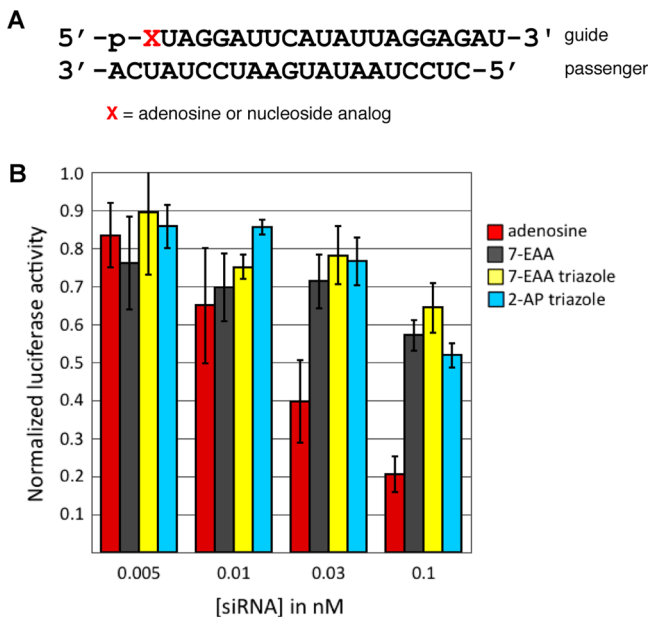


Figure 3. (A) Sequence of siRNA used in this study. X indicates guide strand position 1. All siRNAs were prepared with a 5'-phosphorylated guide strand (p). (B) Knockdown activity of siRNAs with adenosine, 7-EAA, 7-EAA triazole, or 2-AP triazole at the guide strand 5'-end.¹⁸ Activity is reported as the ratio of *Renilla*/firefly luciferase signal at various concentrations of transfected siRNA in HeLa cells.

with the computational results, the 2-AP-triazole inhibited RNAi when present at guide position 1 (Figure 3). The 7-EAA and the 7-EAA-triazole have been tested for their impact on RNAi activity when positioned at the PIK3CB guide strand 5'-end and were re-evaluated here for direct comparison to the 2-AP triazole.¹⁸ Again consistent with the predictions described above, both of these modifications inhibit activity relative to the

parent siRNA with adenosine at position 1 of the guide strand (Figure 3).

While these results suggest computational screening is useful for predicting the effect of different 5'-end modifications, our previously described adenosine derivatives were shown to be poor choices as nucleobase replacements for the siRNA guide strand 5'-end. Others have shown efficient knockdown with an siRNA bearing a nucleobase analogue that is a good mimic of the shape of uridine at this position.¹⁹ However, some loss of activity was reported with a size-expanded adenosine analogue at the 5'-end of the guide strand.²⁰ Considering these observations and the size and shape of the hAgo2 guide strand 5'-end binding pocket, we next evaluated analogues that replace the natural base entirely. We chose to focus on 1,2,3-triazol-4-yl bases since these could be readily prepared from 1-ethynylribose (1-ER) containing precursor RNA and different azides (Figure 4). Such analogues could stack with Tyr 529 in the Ago2 5'-end binding pocket but cannot participate in W/C H-bonding. However, the lack of W/C H-bonding was not considered to be a significant concern for guide strand position 1 because this nucleotide does not directly contact the target RNA strand. We carried out computational screening on 58 different 1,2,3-triazol-4-yl bases, along with the parent 1-ethynylribose, and found compounds that scored well and significantly better than the adenosine derivatives described above (e.g., 1-ER triazoles I and II) (Figure 4; Table 2; Supplementary Figure 1; Supplementary Table 1). The apparent ability of 1-ER and derivative triazoles to fit the hAgo2 guide 5'-end binding site stimulated our synthesis of the required phosphoramidite.

Synthesis of 1-ER Phosphoramidite and Generation of Modified Guide Strands. The DNA type 1-ethynyl 2-deoxyribose phosphoramidite has been synthesized to investigate the effects of various triazole derivatives on the stability of DNA duplexes and triplex-formation.^{21,22} Here we prepared the RNA type 1-ethynyl ribose (1-ER) phosphoramidite to incorporate it into siRNA strands. The skeleton of the β -anomer of 1-ER was synthesized from commercially available sugar **1** using ethyl[trimethylsilyl(ethynyl)]aluminum chloride^{23,24} according to a literature report²⁵ (Scheme 1). Only one isomer was observed in this reaction. The trimethylsilyl and benzoyl groups of **2** were removed with $\text{NH}_4\text{OH}/\text{EtOH}$, and the primary hydroxyl was protected with a dimethoxytrityl group to give compound **3**. The configuration was determined from ^1H NMR NOESY data of compound **3**. Since an NOE between H-1 and H-4 was observed, the isomer was assigned as the β -anomer. Phosphoramidite building block **5** was synthesized by normal *tert*-butyldimethylsilyl protection at the 2-position to give **4** and subsequent phosphorylation.

The 1-ER phosphoramidite building block **5</**

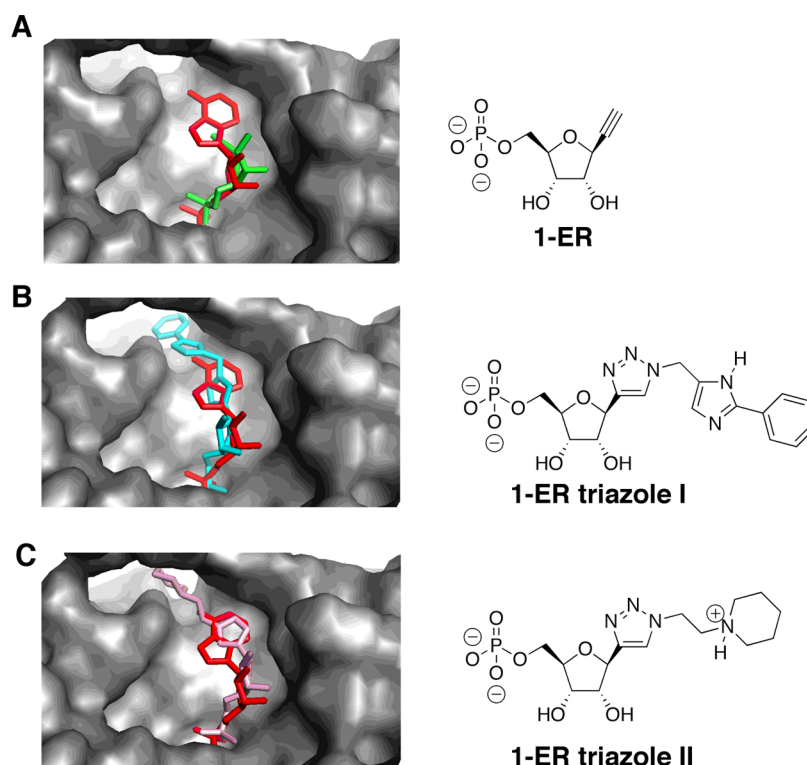


Figure 4. Predicted hAgo2-binding modes for 1-ER derivatives positioned at a guide strand 5'-end. Each analogue is shown in overlay with adenosine (red) in the guide strand 5'-end binding site of hAgo2 shown as a gray surface.¹⁰ (A) 1-ER (green), (B) 1-ER triazole I (blue), (C) 1-ER triazole II (pink).

Table 2

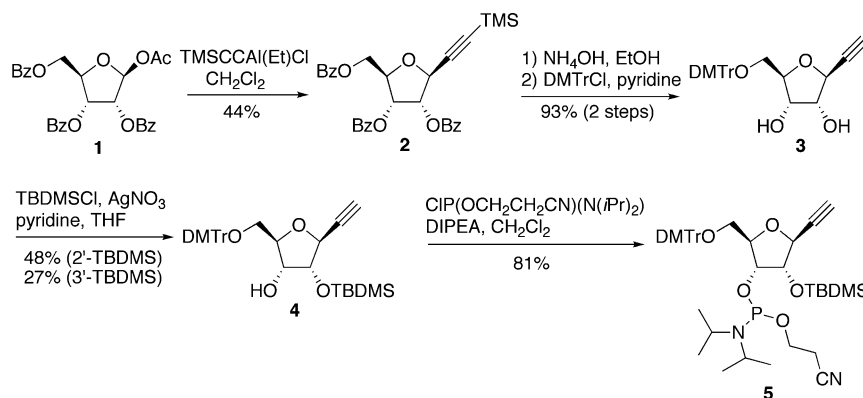
	predicted hAgo2 binding ^a	siRNA activity ^b
adenosine	1.0	++
1-ER	2.1	++
1-ER triazole I	1.2	+++
1-ER triazole II	1.3	+++

^aScores normalized from 1 to 10 from several scoring methods (see Experimental Section). A lower number represents better-predicted binding to guide the strand 5'-end. ^bExperimental siRNA activity determined at 100 pM. +++ = <10% luciferase activity remaining after knockdown; ++ = 10–40%; + = 41–70%; – = >70% (Figure 5).

The reaction was completed in 2–3 h, and the products were purified from the reaction mixture by denaturing PAGE and confirmed by MALDI-MS analysis.

RNA Interference with siRNAs Bearing 1-ER or 1,2,3-Triazol-4-yl Bases. The guide strand of the PIK3CB siRNA was modified with either 1-ER, 1-ER triazole I, or 1-ER triazole II at position 1 (Figure 4). RNAi activity for the modified siRNAs was measured in HeLa cells as described above. As a test of the effect of modification position on activity, we also incorporated these analogues at guide position 12. Less is known about how the base of position 12 interacts with hAgo2 since electron density is lacking for this nucleotide in recently reported hAgo2/guide RNA crystal structures.^{10,11} However, position 12 of the guide strand lies in the middle of the siRNA where pairing to the passenger strand is important for siRNA stability. Furthermore, pairing of target RNA with guide strand nucleotides 2–8 and 12–17 is believed to define the most efficient sites for RNAi, so base structure at guide position 12 would be expected to be important for activity.³¹ For each

Scheme 1



modified siRNA, we carried out a four-point concentration profile in the RNAi assay (Figure 5, Figure 6). Importantly,

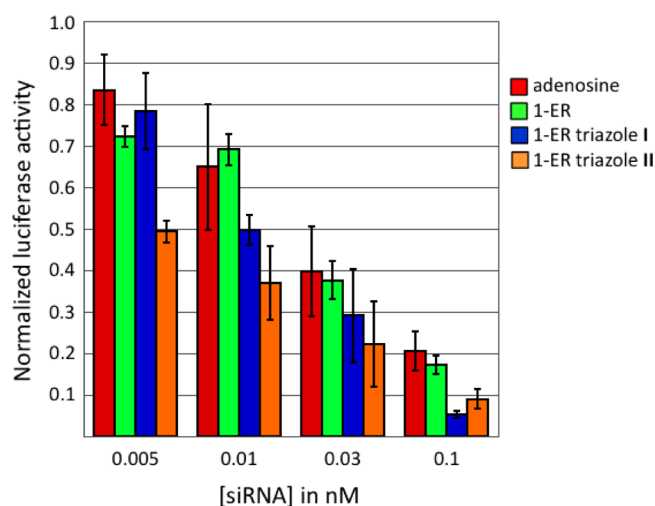


Figure 5. Knockdown activity of siRNAs with adenosine, 1-ER, 1-ER triazole I, or 1-ER triazole II at the guide strand 5'-end. Activity is reported as the ratio of *Renilla*/firefly luciferase signal at various concentrations of transfected siRNA in HeLa cells.

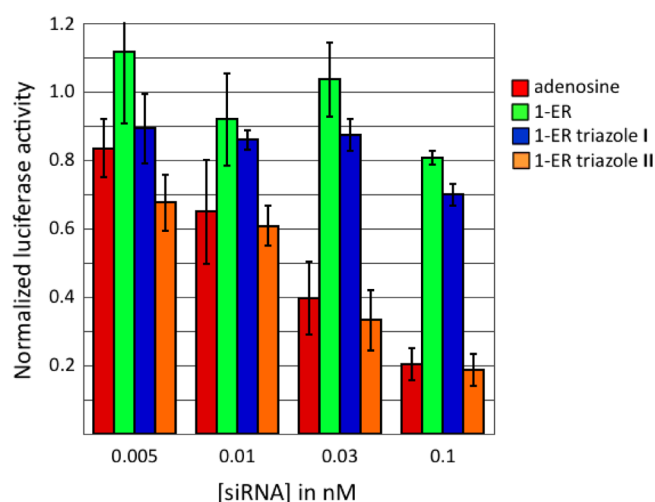


Figure 6. Knockdown activity of siRNAs with adenosine, 1-ER, 1-ER triazole I, or 1-ER triazole II at guide strand position 12. Activity is reported as the ratio of *Renilla*/firefly luciferase signal at various concentrations of transfected siRNA in HeLa cells.

replacement of the adenine base at position 1 with 1-ER, 1-ER triazole I, or 1-ER triazole II was well-tolerated and resulted in siRNAs with potency similar to (1-ER) or exceeding that (1-ER triazoles I and II) of the unmodified duplex (Figure 5). This is in sharp contrast to the purine analogues described above and in good agreement with the computational screening results (Table 2). The docking suggests that 1-ER triazoles I and II are capable of occupying a cleft in the protein adjacent to the guide strand 5'-end binding site and between K525 and P526 of the MID domain and Y815 and L817 of the PIWI domain (Figure 4B,C; Supplementary Figure 2). Crystallization of the complex of hAgo2 bound to a 5'-end modified guide strand will be necessary to test this hypothesis. The activity effects observed are clearly guide strand position-dependent since replacement of the adenine base at guide position 12 with either the 1-ER

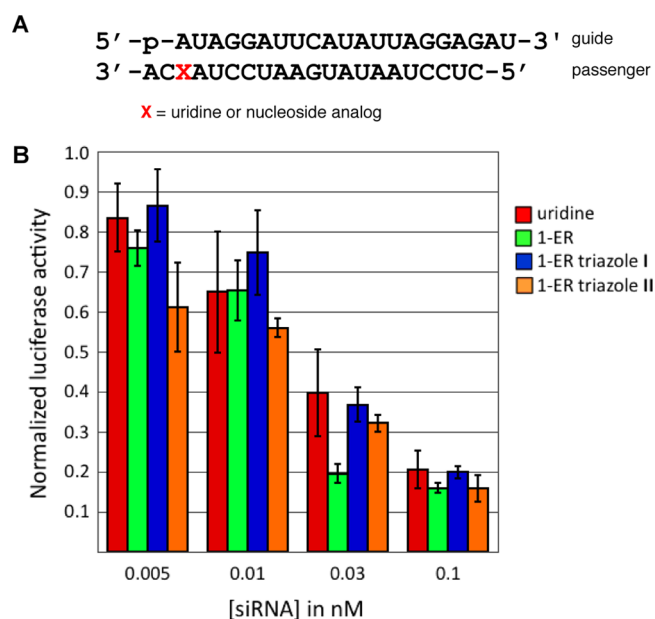
modification or 1-ER triazole I inhibited knockdown (Figure 6).

While the novel modifications may be affecting siRNA potency by modulating guide stand binding to hAgo2 as designed, this is not the only effect that could be responsible for the observed results. The nucleoside analogues could also facilitate duplex unwinding and enhance loading of the desired guide strand into hAgo2.³² This is particularly true for analogues placed near the guide strand 5'-end that are duplex destabilizing (as is the case for 1-ER and the triazoles derived from it, see below, Table 3).³² If this were the source of the

Table 3. Thermal Melting Temperatures of Modified siRNAs

siRNA T_m , °C	guide position 12 modification			
	unmodified	1-ER	1-ER triazole	1-ER triazole II
	64.7 ± 0.4	56.1 ± 0.1	57.0 ± 0.1	56.2 ± 0.3

effects on knockdown efficacy, one would expect to see similar results if the analogue were placed at the same end of the duplex but in the opposite strand. We tested this by preparing siRNAs with 1-ER, 1-ER triazole I, and 1-ER triazole II at position 19 of the passenger strand for the siRNA targeting the PIK3CB sequence (Figure 7A). Interestingly, for the siRNAs



analogue could also be altering strand bias during loading of hAgo2.

We also wished to evaluate the effect these modifications have on the stability of the siRNA to nucleases present in serum since an increase in serum stability could result in increased RNAi activity.³³ However, we found that 1-ER, 1-ER triazole I, and 1-ER triazole II at the guide strand 5'-end caused a slight decrease in serum stability when compared to unmodified siRNA (Supplementary Figure 4). Thus, enhanced RNAi activity arising from these modifications is unlikely to be the result of increasing resistance to nucleases present in serum.

Surprisingly, efficient knockdown is observed with 1-ER triazole II at position 12 (Figure 6). This could be explained by a stabilizing effect on RNA–RNA interactions of a positive charge on the piperidine present in the 1-ER triazole II structure (Figure 4C). We measured thermal melting temperatures (T_M 's) for the modified siRNAs to determine the effect on RNA duplex stability for each of the analogues introduced at guide position 12 (Table 3). All of the modifications made, including 1-ER triazole II, destabilized the siRNA duplex to the same extent. Whereas the unmodified siRNA had a $T_M = 64.7 \pm 0.4$ °C, each of the modified siRNAs had T_M values of 56–57 °C. Therefore, the ability of 1-ER triazole II to function as an effective base replacement at guide position 12 is not due simply to its positive charge stabilizing RNA–RNA interactions. Earlier work showed that 7-EAA and 7-EAA triazole (Figure 2) are effective adenosine replacements at guide position 12.¹⁸ In addition, Li et al. reported that difluorotoluene replacement for uracil at this position resulted in minimal loss in siRNA activity.³⁴ However, Hernandez et al. found that size-expanded nucleobases near the center of an siRNA guide strand (e.g., at positions 9, 10, 11, and 14) caused a substantial reduction in activity.²⁰

These results, along with our observation that 1-ER or 1-ER triazole I at guide position 12 inhibits RNAi, indicate a clear sensitivity to base structure at this position with some nucleobase analogues, but not others, functioning well. The positive effects of certain nucleobase analogues at this position, such as 1-ER triazole II, appear to be largely independent of their effects on duplex stability and likely arise from beneficial interactions with components of the RNAi pathway, perhaps hAgo2 itself. Further study will be necessary to fully define their origin, however.

CONCLUSION

In summary, we developed and validated a method using computational screening for identifying functional 5'-end modifiers for siRNA guide strands. 5'-End modifiers need not be capable of W/C H-bonding but should fit the general shape of the 5'-end binding site in MID/PIWI domains of hAgo2. We devised an efficient synthesis of the 1-ER phosphoramidite and generated RNA bearing this modification. 1-ER serves as a precursor to 1,2,3-triazol-4-yl bases via CuAAC reactions with azides, and specific 1,2,3-triazol-4-yl bases perform well at position 1 of an siRNA guide strand. This stands in contrast to purine derivatives with modified Hoogsteen faces or N2 substituents that inhibit RNAi when placed at the guide strand 5'-end. We also reported the unexpected discovery of a non-W/C H-bonding analogue that performs well at position 12 where H-bonding to target RNA was expected to be important. These studies expand the repertoire of nucleoside analogues useful for siRNAs. In addition, this work sets the stage for future efforts to test our hAgo2 binding hypothesis for 1,2,3-triazol-4-yl bases

with high resolution structures, to evaluate the impact of these novel bases on siRNA off-target effects and their duration of action as well as further computational screening to identify additional high performing guide strand 5'-end modifications.

EXPERIMENTAL SECTION

All reagents were purchased from commercial sources (Sigma–Aldrich or Fisher Scientific) and were used without further purification unless otherwise stated. Reactions were carried out under an atmosphere of dry argon. ¹H, ¹³C, and ³¹P NMR spectra were recorded with Varian VNMR500 or Varian Mercury 300 spectrometers. High-resolution ESI mass spectra were obtained at the University of California, Davis mass spectrometry facility, on an Orbitrap FTMS instrument. Matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectra of oligonucleotides were measured in the positive ion mode, using 6-aza-2-thiothymine as a matrix.

Computational Screening. The overall process of computational screening involved five steps: 1. Library creation, 2. Conformer generation, 3. Receptor creation, 4. Receptor-library docking, and 5. Analysis. All programs mentioned are part of the *OpenEye* Suite of programs.¹⁵ As the approach is aimed at high throughput, molecular dynamics simulations are not part of the docking protocol used.

1. Library Creation. First, a ligand library was created, consisting of molecules thought to be easily accessible from the 7-EAA, 2-AP, or 1-ER structures using commercially accessible precursors from Sigma–Aldrich. This was done to narrow our search to molecules that, if thought to be good candidates after screening, could be quickly generated and incorporated into an oligomer for testing.

2. Conformer Generation. Once this library was compiled, the Omega program was used to generate a list of conformers of each molecule, ranging from <20 for smaller, rigid molecules to >5000 for large, flexible molecules. This required expanding the number of maximum conformers from the program default of 200.

3. Receptor Creation. Starting from published crystallographic data (PDB file 4E11),¹⁰ several similar receptors of various sizes were created, which, along with heavy atom contour surfaces, define how close to the protein surface the heavy atoms in screened ligands can dock. This was accomplished with *make_receptor*. The receptors were created so as to ensure that even the largest molecules in our computational screen would be accommodated in their most extended conformations. Once the heavy atom contours were created, several constraints were applied, to ensure that docking was forced to place the ribose subunit where it would be if it were at the 5'-end of a strand of bound RNA. A constraint was imposed on the ribose 2'-oxygen linkage (modeled in our library as a free hydroxyl). An additional constraint was applied to the ribose 4' oxygen and to the phosphate (see Supplementary Figure 3).

4. Receptor-Library Docking. The library of substrate conformers from step 2 was then docked into the receptors from step 3 using FRED, with several scoring functions available in OpenEye. These scoring methods each have strengths and weaknesses, depending on the system of interest. They utilize various proprietary algorithms that model shape, protein–ligand hydrogen-bonding, implicit solvent–ligand hydrogen bonding, and polarity/electrostatics (see the OpenEye Web site for additional information). Each of these factors are used to give a conformer a score for a specific pose (within the constraints imposed), and once this has been done for all valid poses of all conformers, an overall score and ranking is generated as output, along with atomic coordinates for the corresponding poses.

5. Analysis. Due to the fact that traditional docking programs such as FRED were not specifically designed to model RNA-protein interactions, and the relatively low-level of the calculations involved, we felt further analysis was justified. To account for the fact that we are modeling a flexible protein with a rigid docking method, we analyzed the top 10 poses of the top scoring molecules. We reasoned that, if a molecule is a good fit, the top poses should all score nearly as well as the highest for each candidate. Additionally, these top poses should all bind in a similar way (i.e., molecules

system compared to molecules that show little variance in their top poses). Five sets of scores were created, each differing in the relative importance of the scoring algorithms that contributed to the overall score. For example, the created receptor was scored first with Chemgauss4, the latest scoring function available in OpenEye. Another score was generated with the same receptor, but using the slightly older Chemgauss3 scoring function. Yet another was derived from the Consensus scoring method, which gives a score based on several available scoring functions, as detailed on the OpenEye Web site. Lastly, a method was implemented that utilizes the Consensus method to choose the order of the top poses, which are then actually given a score with Chemgauss3, and then the reverse, being Chemgauss3 pose selection and Consensus scoring. Due to the fact that each of these scoring methods generates scores within their own numerical scale (each scoring function is different), we normalized these scores and averaged them to obtain an overall score. Finally, an ordering of priority for experimental screening was generated for the top scoring molecules based on the criteria of fluxionality described above, and overall rank across all scoring functions. For example, if a molecule was in the top five of four of the five scoring methods and showed little fluxionality, it was deemed an excellent candidate. This last step is more subjective than the normalized scores themselves, as chemical intuition, with knowledge of the limitations of our methods constantly in mind, comes into play.

3,6-Anhydro-1,2-dideoxy-1-(trimethylsilyl)-4,5,7-tri-*O*-benzoyl-*D*-allo-hept-1-ynitol (2). To a solution of trimethylsilylacetylene (1.8 mL, 13 mmol) in anhydrous toluene (3 mL) was added a solution of *n*-BuLi (8.0 mL, 1.6 M solution in hexane, 13 mmol) at 0 °C. After 25 min, to the mixture was added a solution of EtAlCl₂ (13 mL, 1.0 M solution in hexane, 13 mmol). After 30 min, a solution of 1-*O*-acetyl-2,3,5-tri-*O*-benzoyl- β -*D*-ribose **1** (806 mg, 1.6 mmol) in anhydrous CH₂Cl₂ (9 mL) was added, and the mixture was stirred at 0 °C for 2.5 h and at room temperature for 4 h. The reaction mixture was quenched with saturated aqueous NaHCO₃ at 0 °C and filtered through Celite. The filtrate was washed with saturated aqueous NaHCO₃ and dried over Na₂SO₄, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on a silica gel with hexane/EtOAc (19:1→9:1) to give **2** (380 mg, 44%) as a pale yellow oil. ¹H NMR (CDCl₃, 300 MHz, TMS): δ (ppm) 8.08 (d, *J* = 7.8 Hz, 2H), 8.00 (d, *J* = 7.8 Hz, 2H), 7.89 (d, *J* = 7.8 Hz, 2H), 7.60–7.50 (m, 3H), 7.44–7.50 (m, 6H), 5.85–5.78 (m, 2H), 4.96 (d, *J* = 3.6 Hz, 1H), 4.74–4.69 (m, 1H), 4.63–4.56 (m, 2H), 0.17 (s, 9H). ¹³C NMR (CDCl₃, 150 MHz): δ (ppm) 166.4, 165.4, 165.3, 133.7, 133.6, 133.3, 129.9, 128.6, 100.3, 93.9, 79.4, 76.5, 72.9, 72.2, 64.5, –0.22. ESIHRMS (*m/z*): calcd for C₃₁H₃₀O₇Si [M + H]⁺ 543.1834, obsd 543.1839.

3,6-Anhydro-1,2-dideoxy-7-*O*-(4,4'-dimethoxytrityl)-*D*-allo-hept-1-ynitol (3). A solution of 30% NH₄OH was added to a solution of **2** (370 mg, 0.68 mmol) in EtOH, and the mixture was stirred at room temperature for 2 days. The reaction mixture was evaporated under reduced pressure, and the residue was coevaporated with anhydrous acetonitrile and anhydrous pyridine. The dried residue was dissolved in anhydrous pyridine (3 mL). To the solution was added DMTrCl (415 mg, 1.2 mmol), and the mixture was stirred at room temperature for 1 h. The reaction mixture was diluted with EtOAc (100 mL) and washed with water (50 mL) and brine (50 mL). The organic phase was dried over Na₂SO₄, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on a silica gel with hexane/EtOAc (4:1→2:1) containing 0.5% pyridine to give **3** (290 mg, 93%) as a yellow oil. ¹H NMR (CD₂Cl₂, 300 MHz): δ (ppm) 7.47 (d, *J* = 7.8 Hz, 2H), 7.37–7.19 (m, 7H), 6.83 (d, *J* = 9.0 Hz, 4H), 4.47 (dd, *J* = 5.1, 2.1 Hz, 1H), 4.29 (t, *J* = 5.1 Hz, 1H), 4.19 (t, *J* = 5.1 Hz, 1H), 3.99–3.94 (m, 1H), 3.77 (s, 6H), 3.34 (brs, 2H), 3.29 (dd, *J* = 10, 3.9 Hz, 1H), 3.15 (dd, *J* = 10, 4.5 Hz, 1H), 2.60 (d, *J* = 2.1 Hz, 1H). ¹³C NMR (CD₂Cl₂, 150 MHz): δ (ppm) 158.6, 144.9, 136.1, 136.0, 130.3, 130.2, 128.4, 128.0, 126.9, 113.3, 86.5, 83.2, 81.4, 76.8, 75.0, 73.2, 72.8, 64.3, 55.4. ESIHRMS (*m/z*): calcd for C₂₈H₂₈O₆ [M – H][–] 459.1813, obsd 459.1786.

3,6-Anhydro-1,2-dideoxy-7-*O*-(4,4'-Dimethoxytrityl)-4-*O*-(*tert*-butyldimethylsilyl)-*D*-allo-hept-1-ynitol (4). Compound **3** (203 mg, 0.44 mmol) was coevaporated with anhydrous acetonitrile, and the dried residue was dissolved in anhydrous THF (2 mL). To the solution dry pyridine (177 μ L, 2.2 mmol) and AgNO₃ (112 mg, 0.66 mmol) were added. After 5 min, TBDMSCl (119 mg, 0.79 mmol) was added, and the mixture was stirred at room temperature for 2 h. The reaction mixture was diluted with CH₂Cl₂, filtered through Celite, and washed with saturated aqueous NaHCO₃ (20 mL) and brine (20 mL). The organic phase was dried over Na₂SO₄ and concentrated under reduced pressure. The residue was purified by column chromatography on a silica gel with hexane/EtOAc (19:1) containing 0.5% pyridine to give **4** (121 mg, 48%) as a pale yellow foam and 3'-TBDMS (68 mg, 27%) as a pale yellow foam. ¹H NMR (CD₂Cl₂, 300 MHz): δ (ppm) 7.49 (d, *J* = 7.8 Hz, 2H), 7.39–7.22 (m, 7H), 6.84 (d, *J* = 8.7 Hz, 4H), 4.48 (dd, *J* = 6.6, 4.5 Hz, 1H), 4.42 (dd, *J</*

HEPES buffer (pH 7, 12.77 μ L, final concentration 100 mM) was treated with tris-[1-(3-hydroxypropyl)-1H-[1,2,3]triazol-4-yl)methyl]-amine ligand (100 mM solution in H₂O, 1.75 μ L, 175 nmol), either 2-(azidomethyl)-5-phenyl-1H-imidazole (**6**) (100 mM solution in DMSO, 0.35 μ L, 35 nmol) or 1-(2-azidoethyl)-piperidine (50 mM solution in 0.5 M Tris-HCl, pH 8.0, 0.7 μ L, 35 nmol),³⁶ sodium ascorbate (100 mM solution in H₂O, 1.75 μ L, 175 nmol), and CuSO₄ (100 mM solution in H₂O, 0.88 μ L, 88 nmol). The solution was incubated for 2–3 h at room temperature. The reaction mixture was then diluted with PAGE loading buffer (80% formamide containing 10 mM EDTA) and PAGE-purified (19% denaturing, 20 W, 6–8 h). After electrophoresis, the RNA bands were visualized by UV shadowing (254 nm light, F254 TLC plate as a backing) and extracted from the gel via the crush and soak method at 4 °C overnight in 0.5 M NH₄OAc containing 0.1 mM EDTA. Polyacrylamide particles were removed using a Centrux filter (0.2 μ m), and the oligonucleotide solution was desalted using C18 Sep-Pak cartridges (Waters), eluting with 1:1 CH₃CN/H₂O or precipitated with 3:1:0.1 ethanol/crush and soak oligonucleotide solution/3 M sodium acetate followed by 70% ethanol wash. 1-ER containing RNAs were purified by gel electrophoresis and isolated as described above for the triazole-containing RNAs.

RNA Quantification and Mass Spectrometry Analysis. The oligonucleotide solutions or precipitated pellets were lyophilized to dryness, resuspended in nuclease-free water, and quantified by absorbance measurements at 260 nm. Click reaction yields for guide position 1 ($84 \pm 2\%$ for 1-ER triazole I and $81 \pm 2\%$ for 1-ER triazole II, values averaged over three trials $\pm$ standard deviation) were determined by comparing the side-by-side purified yield of unclicked 1-ER oligonucleotides from the crude mixture to the click reactions on the same crude RNA. Identity of the oligonucleotides containing either 1-ER, triazole I, or triazole II modifications was confirmed by MALDI-TOF mass spectroscopy using a saturated solution of 6-aza-2-thiothymine in 0.1 M aqueous dibasic ammonium citrate as a matrix. Mass spectra were recorded in the positive ionization mode and calibrated to an internal DNA standard of 6740.4 Da. List of mass values, $[M + H]^+$, (G = guide, P = passenger, number indicates position of modification), for all 1-ER modified siRNAs: G1: calcd 6718.0, obsd 6719.0; G12: calcd 6718.0, obsd 6719.2; P19: calcd 6477.0, obsd 6477.3. For all 1-ER triazole I modified siRNAs: G1: calcd 6917.2, obsd 6717.7; G12: calcd 6917.2, obsd 6920.2; P19: calcd 6676.2, obsd 6676.4. For all 1-ER triazole II modified siRNAs: G1: calcd 6872.2, obsd 6873.1; G12: calcd 6872.2, obsd 6872.5; P19: calcd 6631.2, obsd 6630.1.

Melting Temperature Analysis. The thermal stability of siRNA duplexes containing either adenosine, 1-ER, 1-ER triazole I, or 1-ER triazole II at guide position 12 were analyzed in a similar fashion to our previously described methods.^{16,18} siRNA duplexes were formed by hybridizing 462.5 pmol of complementary strands in 925 μ L of TE buffer (10 mM Tris-HCl, pH 7.8, 0.1 mM EDTA, and 100 mM NaCl). The values reported in Table 3 are an average of three thermal melting experiments (152.5 pmol, 305 μ L per experiment), with an experimental temperature range of 30–80 °C. The values in Table 3 indicate the average melting temperature $\pm$ standard deviation.

Cell Culture. HeLa cells (ATCC) were grown at 37 °C in humidified 5% CO₂ in Dulbecco's modified Eagle's medium (DMEM, GIBCO) supplemented with 10% fetal bovine serum (FBS, GIBCO) and 1% antibiotic-antimycotic (100 $\times$ Anti-Anti, GIBCO). The cells were maintained in exponential growth.

Transfection and RNAi Activity Assay. For the dual-luciferase RNA interference assay, HeLa cells were grown to 80–90% confluence then detached using Accutase (Innovative Cell Technologies, without calcium or magnesium). Reverse transfection of the HeLa cells was accomplished using siPORT NeoFX (Ambion) in Opti-MEM media (GIBCO) to deliver the psiCHECK-2-PIK3CB plasmid and siRNAs as previously described.¹⁸ siRNA duplex hybridization was accomplished by combining equal amounts of purified passenger and native or modified guide strands to a final concentration of 5 μ M in 10 mM Tris-HCl, 50 mM KCl, pH 7.5. The samples were heated at 95 °C for 5 min followed by slow cooling to room temperature over a period of

approximately 2 h. All siRNAs referenced in Figures 3, 5, and 6 were tested side-by-side in the same 96-well plate assay. Three individual assays were averaged to give the values and standard deviations plotted in those figures.

■ ASSOCIATED CONTENT

● Supporting Information

Structures and computational screening scores for analogues evaluated, supplementary figures with docking constraints and putative 1,2,3-triazol-4-yl binding site in hAgo2, NMR spectra for compounds 2–5. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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