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Reversible Photocycloaddition of 8-Pyrenylvinylguanine for Photoreactive Serinol Nucleic Acid (SNA)

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Photoresponsive nanomachines are attractive components of functional nanodevices and nanosystems. To develop new photoresponsive nucleic acid-based nanomachines, we conjugated 8-pyrenylvinyl guanine (^{PV}G) to a serinol linker and incorporated it into acyclic xeno nucleic acid, serinol nucleic acid (SNA). The two ^{PV}G residues incorporated into SNA underwent interstrand photocycloaddition upon 447 nm light irradiation in the duplex state, whereas previously reported 8-pyrenylvinyl adenine (^{PV}A) formed both intrastrand and interstrand photodimers. The ^{PV}G photodimer was converted to monomers by irradiation with 350 nm light. This photoreaction enabled reversible photoregulation of formation and dissociation of SNA/RNA duplex, although some byproducts were generated due to slower photoreaction of P^VGs than of ^{PV}As. In contrast, when a single ^{PV}G was incorporated into SNA, the interstrand photocycloaddition and cycloreversion were remarkably fast and effective in the single-stranded state. We utilized this to demonstrate a photocaging system that achieves one-way photoswitching of hybridization ability. The powerful photocycloaddition properties of ^{PV}G-SNA are expected to find applications in new photoresponsive nanodevices and nanosystems.

Introduction

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DNA nanotechnology, such as DNA nanostructures,¹ DNA computings,² DNA-based nanomachines,³ and molecular robotics⁴ offers an attractive platform for realizing highly functional nanodevices and biological tools. Chemical modifications on nucleobases and additional incorporation of functional molecules into the DNA strands facilitate the development of stimuli-responsive systems triggered by metal ions,⁵ pH,⁶ and photoirradiation,⁷ providing powerful systems for the design of novel nanomachines based on nucleic acids.

To overcome the limitations of conventional DNA nanotechnology, it is essential to utilize xeno nucleic acids (XNAs)⁸ with modified backbone structures that exhibit high resistance to enzymatic and chemical degradation and unique hybridization properties distinct from those of natural nucleic acids. In fact, several examples of nanotechnology utilizing XNA have been reported, demonstrating greater functionality than those utilizing DNA.⁹ We previously reported a reversible photoregulation of duplex formation and dissociation of serinol nucleic acid (SNA)¹⁰, acyclic XNA, by introducing two 8-pyrenylvinyladenine (^{PV}A) residues.¹¹ When blue light is irradiated onto the duplex composed of SNA containing two ^{PV}A residues and complementary RNA, [2+2] photocycloaddition reaction occurs between two ^{PV}A residues, causing the duplex to dissociate into single strands. On the other hand, upon UV

light irradiation, the ^{PV}A photodimer undergoes photocycloreversion to regenerate monomeric ^{PV}A, enabling the re-formation of the duplex. This system shows promise for the development of photon-driven nanomachines and photocontrollable biological tools that use XNA as a building block. However, these strategies are limited to photoresponsive modification of adenine base, requiring two adjacent adenines to design photocontrollable system, which limits the applicable sequences.



Fig. 1 (A) Schematic of previous work on the SNA strand modified with $^{\rm PV}A$ residues and (B) this work on the SNA strand modified with $^{\rm PV}G$ residues.

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We report on the photoregulation of SNA containing a photoresponsive nucleobase, 8-pyrenylvinylguanine (^{PV}G). The synthesis of ^{PV}G incorporated into deoxyribose has already been reported.¹² Ogasawara demonstrated that ^{PV}G-modified DNA can be used as a photoswitch for G4 formation via the *trans-cis* isomerization of ^{PV}G, which was successfully applied to photoregulation of the transcription system.¹³ We hypothesized that two adjacent ^{PV}G residues conjugated with a flexible SNA strand are more likely to cause a photocycloaddition than *cis*-isomerization as in the case of ^{PV}A, which should lead to a photoregulation system different from the DNA-^{PV}G system based on a *trans-cis* isomerization. Therefore, we synthesized a novel ^{PV}G-SNA phosphoramidite monomer and incorporated it into the SNA strand, evaluated the photoreactivity of the ^{PV}G residue, and compared it with the ^{PV}A modification.

Results and discussion

Synthetic procedures for the ^{PV}G monomer.

First, we prepared a phosphoramidite monomer of ^{PV}G-SNA according to Fig. 2. The N-9 position of 2-amino-6-chloropurine was converted to the benzyl ester form of compound 1, and then the 6-position was substituted with oxygen to construct the guanine structure 3. The formamidine protection on the 2amino group resulted in an unexpected ester exchange from benzyl ester to ethyl ester 4, but we continued the synthesis. After 8-bromination by NBS, the Cul/CsF-mediated Stille reaction¹⁴ was carried out to obtain the 8-vinylguanine skeleton 6. The Heck reaction with 1-bromopyrene followed by hydrolysis gave 2-protected-^{PV}G 8. After coupling with DMTr-Lserinol, the desired phosphoramidite monomer 10 was successfully obtained using conventional phosphoramidite chemistry. The SNA strand with two ${}^{\mbox{PV}}\mbox{G}$ residues was synthesized using a DNA synthesizer, with each SNA monomer prepared as reported previously.¹⁰



Fig. 2 Synthesis of phosphoramidite monomer of ^{PV}G-SNA **10**. (i) Benzyl bromoacetate, DIPEA, DMF, 0 °C to r.t., o.n., (ii) K₂CO₃, DABCO, Benzyl alcohol, 80 °C, 16 h, (iii) TFA, DCM, r.t., 1 h, (iv) *N*,*N*-Dimethylformamide diethyl acetal, EtOH, 50 °C, 16 h, (v) NBS, DMF, r.t., 2 h, (vi) Pd(PPh₃)₄, Tributyl(vinyl)tin, Cul, CsF, DMF, 55 °C, 4 h, (vii) 1-Bromopyrene, Pd(Ac)₂,PPh₃, NEt₃, DMF, 110 °C, 3 h, (viii) NaOH, 1,4-Dioxane, MeOH, EtOH, H₂O, r.t., 20 min, (ix) DMTr-L-serinol, DMT-MM, NEt₃, DMF, 0 °C for 20 min to r.t. for 1 h, (x) 2-Cyanoethyl *N*,*N*-diisopropylchlorophosphoramidite, NEt₃, CH₂Cl₂, 0 °C, 30 min.

Comparison of the photoreactivity between ^{PV}G and ^{PV}A introduced into the SNA strand.

The melting temperature of the SNA-2^{PV}G/RNA-2C duplex was 40.4 °C (Fig. 3A), which is slightly higher than that of SNA-2^{PV}A/RNA-2U (31.0 °C, Fig. S1). The stabilization of the duplex is probably caused by strong stacking interaction of PVGs, as will be explained later. This result shows that SNA-2^{PV}G and RNA-2C form a duplex at room temperature, suggesting that the PVG substitution does not suppress duplex formation. Subsequently, the photoreactivity of ^{PV}G in the SNA strand was evaluated (Fig. 3). Upon irradiation with 447 nm light to SNA-2^{PV}G/RNA-2C duplex at 20 °C, the absorption band at around 400 nm decreased with the irradiation time, followed by the appearance of a new absorption band at around 330-360 nm, which was attributed to the pyrene moiety (Fig. 3B). The isosbestic points were 329 and 361 nm. This behavior is similar to the photocycloaddition of PVA residues in SNA-2PVA/RNA-2U duplex, indicating that ^{PV}G residues also form a photodimer via photocycloaddition rather than cis-isomerization under the same conditions (Fig. S2).12 When the photodimer of ${}^{\rm PV}\!G$ was irradiated with 350 nm UV light, the absorption bands of the monomeric ^{PV}G were restored (Fig. 3C), indicating a

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cycloreversion reaction similar to that of ^{PV}A. We confirmed that pyrimidine dimers are not generated even when exposed to 350 nm light for 30 minutes under the present conditions (Fig. S12). However, the kinetics of the photocycloaddition of **SNA-** $2^{PV}G/RNA-2C$ were rather slow compared to **SNA-** $2^{PV}A/RNA-$ **2U**: the half-life of the photodimerization of ^{PV}As was about 10 s (Fig. 3D), while that of ^{PV}Gs under the same conditions was over 10 min. Furthermore, the efficiency of cycloreversion also differed: 79% of ^{PV}A photodimer was converted to monomers after 10 min of UV irradiation, whereas only 67% of ^{PV}G photodimer was converted to monomers even after 30 min of UV irradiation (Fig. 3E).



Fig. 3 (A) Chemical structures of SNA units, sequences used in this study, and schematics of the irradiation experiment. (B) Absorption spectra of **SNA-2^{PV}G/RNA-2C** at the indicated times of irradiation with 447 nm light. (C) Absorption spectra of **SNA-2^{PV}G/RNA-2C** after 1 h of irradiation with 447 nm and after irradiation for the indicated times with 350 nm light. Irradiation was performed at 20 °C. (D)(E) The ratio of monomeric ^{PV}G in **SNA-2^{PV}G/RNA-2C** (purple circles) and that of monomeric ^{PV}A in **SNA-2^{PV}A/RNA-2U** (blue triangles, Fig. S2 for UV spectra) as a function of irradiation time (D) at 447 nm light and (E) at 350 nm light. Before irradiation with 350 nm light, ^{PV}G and ^{PV}A were crosslinked by irradiation with 447 nm light for 1 h and 10 min, respectively. The ratio of the monomers was calculated from the absorbance at 400 nm.

To understand the reason for the different behavior of PVG and PVA, we performed denaturing PAGE analysis before and white light irradiation (Fig. 4A,B). To visualize the SNA strand, the entire gel was irradiated with 350 nm light after electrophoresis to restore the fluorescence of the monomeric ${}^{PV}G/{}^{PV}A$ and detected by an imaging analyzer with a blue laser. Note that both ^{PV}G and ^{PV}A are emissive by blue light excitation. In the case of SNA-2^{PV}A/RNA-2U, irradiation with 447 nm resulted in the appearance of two bands and the disappearance of the original band. The lower and the upper bands are assigned to the intrastrand and the interstrand crosslinking products, respectively (Fig. 4A). Under the present conditions employed, PVAs were able to crosslink not only intrastrand but also interstrand. This fact suggests that dimer of the SNA-2^{PV}A/RNA2U duplexes is temporarily formed as an intermediate via association between PVAs in the duplexes. Subsequent irradiation with 350 nm significantly reduced these bands and regenerated the original band of SNA-2^{PV}A due to cycloreversion. In contrast, irradiation of SNA-2^{PV}G/RNA-2C with 447 nm resulted in two additional bands in addition to the remaining unreacted SNA-2^{PV}G strand (Fig. 4B, left). The upper high-intensity band is attributed to the product of interstrand crosslinking because it showed significantly reduced mobility compared to intrastrand-crosslinked SNA-2^{PV}A. The lower band originated from unidentified by-products. Importantly, no band could be attributed to the intrastrand ^{PV}G-dimer, suggesting that intrastrand crosslinking of ^{PV}G was strongly suppressed. Upon irradiation with 350 nm light, the crosslinking product was converted to SNA-2^{PV}G with ^{PV}G monomers, whereas the byproducts remained. Unlike SNA-2PVA/RNA2U, the SNA-2^{PV}G/RNA-2C duplex may facilitate a static stacking interaction between the interstrand ^{PV}Gs to form a dimer of the duplexes prior to photocrosslinking (Fig. 4C). In this case, it is difficult for PVG to adopt the necessary conformation for crosslinking, due to the rigid structure of the duplexes and the electrostatic repulsion between them.

This hypothesis was supported by the results of the irradiation experiments in the absence of complementary RNA. Irradiation with 447 nm light for single-stranded SNA-2^{PV}G showed significantly faster photocrosslinking than in the duplex state (Fig. 4D and S3). The reaction half-life for photocycloaddition was approximately 1 min, which is about 10 times faster than that of the duplex state. The efficiency of the cycloreversion reaction for SNA-2^{PV}G was also improved: approximately 80% of the crosslinked ^{PV}G was converted to the monomer by irradiation with 350 nm for 20 min. Denaturing PAGE revealed that the main product of photoirradiation on single-stranded SNA-2^{PV}G was also only interstrand crosslinking product and that the formation of by-products was inhibited (Fig. 4B, right). The rapid photocycloaddition of single-stranded SNA-2^{PV}G can be explained by the fact that the single-stranded state facilitates the interaction between PVGs on different strands due to the smaller electrostatic repulsion, and that the flexibility of the single strand allows dimerized structure of consecutive PVGs suitable for photocrosslinking reaction compared to the rigid duplex state. As a result, the efficiency of cycloreversion was also enhanced due to the relative suppression of the irreversible

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side reaction during irradiation with 447 nm light. The concentration dependence of the photocrosslinking kinetics provided further evidence for the interstrand reaction of ^{PV}Gs. The crosslinking rate of 1 μ M **SNA-2^{PV}G** with 447 nm light was found to be slower than that of 5 μ M **SNA-2^{PV}G** (Fig. S3 and S4). We conclude that two ^{PV}Gs introduced into the SNA strand undergo photocycloaddition reactions in an interstrand manner.



Fig. 4 (A)(B) Gel-shift assay with denaturing PAGE. (A) **SNA-2**^{PV}**A/RNA-2U** duplex, (B, left) **SNA-2**^{PV}**G/RNA-2C** duplex, and (B, right) **SNA-2**^{PV}**G** single strand. The fluorescence of ^{PV}G and ^{PV}A was detected by excitation at 473 nm laser and long-path filter (510 nm). (C) Illustration of different mechanisms of photocycloaddition between ^{PV}G and ^{PV}A. (D) The ratio of monomeric ^{PV}G in **SNA-2**^{PV}**G** single strand at irradiation with 447 nm (red circles) and 350 nm (blue squares) (E) Melting profile of **SNA-2**^{PV}**G/RNA-2C** before (black line) and after irradiation with 447 nm light (purple line) and 350 nm light (blue line).

Although the reaction mechanism was different from what we expected, we hypothesized that the photoreaction of ^{PV}Gs would enable photoregulation of duplex formation. In **SNA-**2^{PV}A/RNA-2U, which induces both intrastrand and interstrand crosslinks, clear photoregulation of duplex formation and dissociation was observed, which was reversible and consistent with our previous results (Fig. S5).¹¹ Irradiation of the **SNA-**2^{PV}G/RNA-2C</sup> duplex with 447 nm light at 20 °C led to the complete disappearance of the sigmoidal melting curve, indicating effective dissociation of the duplex by interstrand

photocycloaddition of ^{PV}Gs (Fig. 4E, purple \sqrt{line})_{ticle}Upon irradiation of the photoadduct with 350 RAI light, We signal a curve partially recovered, supporting the reformation of the duplex (Fig. 4E, blue line). However, ^{PV}G-SNA showed poor reversibility of the photoreaction (Fig. S6), which is likely due to an unidentified by-product during the irradiation.

Although the precise mechanism remains unclear, it is postulated that ${}^{PV}G$ has a favorable orientation in photocycloaddition reactions, enabling it to photocrosslink solely via interstrand ${}^{PV}G$ dimerization (Fig 4C). In addition, the strong stacking interaction between ${}^{PV}Gs$ facilitates interstrand association during photoreaction (Fig. S7). The formation of byproducts could be the reason for the low cycloreversion efficiency of the ${}^{PV}G$ system compared to the ${}^{PV}A$ system.

Single ^{PV}G incorporated into the SNA strand for effective interstrand photocrosslinking.

By focusing on the strong stacking property and selective interstrand photocycloaddition between ^{PV}Gs, we hypothesized that the SNA strand carrying a single ^{PV}G should undergo interstrand crosslinking and form a 4-arm branched SNA (Fig. 5A). We therefore prepared SNA-1^{PV}G, which carried only a single ^{PV}G residue in the SNA strand, and measured the absorption spectra after photoirradiation (Fig. 5B,C). Upon irradiation with 447 nm light on single-stranded SNA-1^{PV}G, a slight blue shift of the absorption band was observed as an intermediate product, which was attributed to the cis-isomer of ^{PV}G . It subsequently disappeared with the appearance of the pyrene band (Fig. 5B). The isosbestic point was not observed. Presumably, ^{PV}G in **SNA-1**^{PV}G promotes rapid trans-to-cis isomerization of ^{PV}G, and *cis*-^{PV}G is also excited and reverted to the trans form, causing photocrosslinking by 447 nm light. SNA-2^{PV}G involving consecutive ^{PV}Gs probably suppresses cis isomerization through strong stacking interaction between ^{PV}Gs. The reaction rate of photocycloaddition for single stranded SNA-1^{PV}G was as fast as that for single-stranded SNA-2^{PV}G (Fig. 5D). Although *cis*-isomerization competed with photocrosslinking, the photocycloaddition reaction was remarkably fast. Fujimoto et al. reported that [2+2] photocycloaddition of 3-cyanovinylcarbazole (CNVK) occurred only with the trans isomer.¹⁵ In addition, the [2+2] photocycloaddition reaction is generally selective for transisomers.¹⁶ We therefore hypothesize that *cis* ^{PV}G may also need to revert to the trans form prior to photocycloaddition (Fig. S8). However, since some stilbene derivatives have been reported to undergo photodimerization between trans- and cisisomer,^{7h,17} it cannot be completely ruled out that the cis-PVG may undergo a [2+2] photocycloaddition reaction. Upon irradiation of the photocrosslinked SNA-1PVG with 350 nm, the initial absorption band of ^{PV}G was rapidly restored, demonstrating that SNA-1^{PV}G underwent highly effective cycloreversion compared to SNA-2^{PV}G (Fig. 5C,D). Repeated irradiation experiments revealed that an effective photoreaction occurred for at least 5 cycles without severe photobleaching (Fig. S9). Denaturing PAGE also confirmed the selective formation of the interstrand crosslinking product and

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the effective back reaction (Fig. 5E). The concentration dependence of the photoreaction confirmed an interstrand crosslinking reaction between $^{\rm PV}Gs$: the difference in kinetics between 1 μM and 5 μM was approximately fivefold (Fig. S10, S11).



Fig. 5 (A) Schematic of the photoreaction of single-stranded SNA-1^{PV}G by light irradiation. (B) Absorption spectra of single-stranded SNA-1^{PV}G at the indicated times of irradiation with 447 nm light. (C) Absorption spectra of single-stranded SNA-1^{PV}G after 1 h of irradiation with 447 nm and after irradiation for the indicated times with 350 nm light. Irradiation was performed at 20 °C. (D) The ratio of monomeric ^{PV}G in SNA-1^{PV}G single strands after irradiation at 447 nm (red circles) and 350 nm (blue squares). (E) Gel shift assay using denaturing PAGE before and after the irradiation of light to SNA-1^{PV}G single strand.

Photocaging strategy of ${}^{\rm PV}\mbox{G-tethered SNA}$ for the photoactivation of hybridization with RNA.

We demonstrated the one-way photoswitching of the hybridization ability of **SNA-1**^{PV}**G** by photocycloreversion using the pre-dimerized strand (Fig. 6A). The Single-stranded **SNA-1**^{PV}**G** was irradiated with 447 nm light for 20 min to ensure complete photodimerization. Complementary RNA (**RNA-2C**) was then added, and the melting temperature was measured. The melting profile showed no transition, clearly indicating that the photocrosslinked **SNA-1**^{PV}**G** had lost its hybridization ability (Fig. 6B). After irradiating this sample with 350 nm light for 10 min, a sigmoidal curve was observed due to duplex formation. Thus, we confirmed that the photocaging strategy with precrosslinked **SNA-1**^{PV}**G** is feasible. We also confirmed that single modification with ^{PV}A is applicable to photocaging strategy (Fig. S13).





Experimental

Materials

Reagents for the synthesis of the ^{PV}G phosphoramidite monomer were purchased from Tokyo Kasei Co., Ltd., Wako Pure Chemical Industries, Ltd., and Aldrich. The reagents for oligomer synthesis and Poly-Pak II cartridges were purchased from Glen Research. The column for HPLC purification was purchased from Kanto Chemical Co., Ltd. RNA strands were purchased from Hokkaido System Science Co., Ltd.

Photoirradiation

A xenon light source (MAX-301, Asahi Spectra) equipped with interference filters centered at 350 nm and 447 nm lightemitting diodes (LEDs; CCS) was used for photoirradiation. The sample solution was added to a cuvette, and the temperature of the light irradiation was controlled using a programable temperature controller. All photoirradiation experiments were conducted at 20 °C.

Spectroscopic measurements

UV/Vis spectra were measured using a JASCO model V-730 spectrometer equipped with a programable temperature controller; 10-mm quartz cells were used. The sample solutions contained 100 mM NaCl and 10 mM phosphate buffer (pH 7.0). The duplex concentration was 5.0 μ M, and the experiments were performed in a 10-mm quartz cell.

Melting-temperature measurements

The melting curves of all nucleic acid duplexes were obtained using a JASCO model V-730 spectrometer equipped with a programable temperature controller by measuring the change in absorbance at 260 nm versus temperature. The melting curves for heating and cooling were measured, and the average maximum value of their first derivatives was determined to be the melting temperature (T_m). The solution conditions were (unless otherwise noted) 100 mM NaCl, 10 mM phosphate buffer (pH 7.0), 5.0 μ M oligonucleotide.

Denaturing PAGE

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Samples were loaded into the wells of a denaturing gel (20% acrylamide, 8 M urea, and 1×TBE). Electrophoresis was performed at 4 W at r.t. for 2 h. The gels were irradiated with 350 nm light after electrophoresis to recover the fluorescence of the monomeric ^{PV}G and ^{PV}A. The gels were analyzed using a Typhoon FLA 9500 with a 473 nm laser and LPB filter (510 nm).

Synthesis of compound 1

A suspended solution of 2-amino-6-chloropurine (5.00 g, 29.5 mmol) and *N*,*N*-diisopropylethylamine (5.53 mL, 32.5 mmol) in dry DMF was stirred in an ice bath for 10 min. Then, benzyl bromoacetate (5.55 g, 35.4 mmol) was added to the solution. After stirring 1 h on an ice bath, stir again at an ambient temperature overnight. After evaporation, the residue was suspended in EtOH and filtered to collect the solid. The solid was washed with diethyl ether and dried under vacuo to give **1** (6.77 g, 73%). ¹H-NMR [DMSO-d₆, 500 MHz] δ = 8.12 (s, 1H), 7.41-7.32 (m, 5H), 7.00 (s, 1H), 5.17 (s, 1H), 5.06 (s, 1H). ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 168.1, 160.4, 154.7, 149.9, 143.9, 135.8, 129.0, 128.8, 128.6, 123.4, 67.2, 44.5. HRMS (FAB) Calcd for C₁₄H₁₂ClN₅O₂ (M+H⁺): 318.0758. Found 318.0761.

Synthesis of compound 2

Under N₂ atmosphere, benzyl alcohol (35 mL) was added to Compound **1** (6.79 g, 21.4 mmol), K₂CO₃ (3.00 g, 21.4 mmol), and 1,4-Diazabicyclo[2.2.2]octane (0.240 g, 2.14 mmol). The mixture was stirred at 80 °C for 16 h. After cooling to room temperature, chloroform was added to the mixture and washed with water. The water layer was washed with chloroform, and the combined organic layer was subsequently washed with brine. The collected organic layer was evaporated. The residue was suspended with diethyl ether and then filtered to give **2** as a solid (5.21 g, 63%). ¹H-NMR [DMSO-d₆, 500 MHz] δ = 7.86 (s, 1H), 7.52-7.31 (m, 10H), 6.53 (s, 1H), 5.50 (s, 1H), 5.18 (s, 2H), 5.01 (s, 2H) . ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 168.4, 160.5, 160.3, 155.1, 140.6, 137.1, 135.9, 129.0, 128.9, 128.7, 128.54, 128.52, 113.7, 67.4, 67.1, 44.3. HRMS (FAB) Calcd for C₂₁H₁₉N₅O₃ (M+H⁺): 390.1566. Found 390.1560.

Synthesis of compound 3

Compound **2** (5.21 g, 13.4 mmol) was dissolved in a trifluoroacetate: $CH_2Cl_2 = 1:1$ solution and stirred for 1 h at room temperature. After brief evaporation, the residue was co-evaporated with acetonitrile three times. The residue was suspended in diethyl ether and then filtered to give **3** as a solid (4.91 g, >100%).¹H-NMR [DMSO-d₆, 500 MHz] δ = 10.88 (br, 1H), 7.88 (s, 1H), 7.40-7.32 (m, 5H), 6.72 (br, 1H), 5.19 (s, 1H), 4.97 (s, 1H). ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 167.8, 156.3, 154.3, 151.3, 137.9, 135.4, 128.6, 128.4, 128.1, 67.1, 44.5. HRMS (FAB) Calcd for C₁₄H₁₃N₅O₃ (M+H⁺): 300.1096 Found 300.1094.

Synthesis of compound 4

N,*N*-dimethylformamide diethyl acetal (40 mL, 164 mmol) was added to compound **3** (4.91 g, 13.4 mmol), which was suspended in EtOH. The mixture was stirred for 16 h at 50 °C. After evaporation, it was re-precipitated by the addition of a small amount of cold ethyl acetate, then cooled on an ice bath.

The precipitate was collected by filtration and dried in Nacua to give **4** (2.67 g, 68%).¹H-NMR [DMSO-d₆, 500 MH2F8/E5PE29(bf, 1H), 8.52 (s, 1H), 7.79 (s, 1H), 4.95 (s, 2H), 4.17 (q, 2H), 3.13 (s, 3H), 3.02 (s, 3H), 1.22 (t, 3H). ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 168.5, 158.5, 158.1, 157.8, 150.6, 139.4, 119.3, 61.8, 44.2, 41.1, 35.1, 14.5. HRMS (FAB) Calcd for C₁₂H₁₆N₆O₃ (M+H⁺) 293.1362. Found 293.1343.

Synthesis of compound 5

Compound 4 (2.67 g, 9.13 mmol) was dissolved in dry DMF. N-Bromosuccinimide (1.96 g, 11.0 mmol) in dry DMF was added dropwise to the solution. The mixture was stirred for 2 h under N₂ atmosphere. The reaction was quenched by the addition of 5 % $Na_2S_2O_4$ aq. (dry DMF : 5 % $Na_2S_2O_4$ aq.=1:1) on an ice bath. After stirring for 5 min, the solution was extracted with chloroform and $Na_2S_2O_4$, and the organic layer was collected. The water layer was washed with chloroform. The combined organic layer was evaporated. The residue was suspended in small amounts of chloroform and large amounts of diethyl ether. The precipitate was collected by filtration and dried in vacuo, yielding **5** (3.11 g, 92%).¹H-NMR [DMSO-d₆, 500 MHz] δ = 11.46 (br, 1H), 8.59 (s, 1H), 4.91 (s, 2H), 4.20 (q, 2H), 3.15 (s, 3H), 3.03 (s, 3H), 1.23 (t, 3H). $^{13}\text{C-NMR}$ [DMSO-d_6, 125 MHz] δ = 167.6, 158.9, 158.2, 156.9, 151.9, 123.0, 119.6, 62.2, 44.7, 41.2, 35.2, 14.5. HRMS (FAB) Calcd for C₁₂H₁₅BrN₆O₃ (M+H⁺): 371.0467 Found 371.0482.

Synthesis of compound 6

(700 Compound 5 1.89 mg, mmol), tetrakis(triphenylphosphine)palladium(0) (320 mg, 0.28 mmol), cupper iodide (70 mg, 0.38 mmol), and cesium fluoride (570 mg, 3.78 mmol) were suspended in dry DMF. The solution was then added tributyl(vinyl)tin (1.1 mL, 3.78 mmol) and stirred at 55 °C for 4 h. The solution was filtered using Cerite to remove the insoluble solid. After evaporation, the residue was purified by silica gel column chromatography (CHCl₃:MeOH = 20:1), yielding **6** (280 mg, 47%). ¹H-NMR [DMSO-d₆, 500 MHz] δ =11.33 (s, 1H), 8.57 (s, 1H), 6.71 (m, H), 6.17 (s, 1H), 5.47 (d, 1H), 5.06 (s, 2H), 4.17 (q, wH), 3.14 (s, 1H), 3.03 (s, 1H), 1.21 (t, 3H). ¹³C-NMR $[DMSO-d_6, 125 MHz] \delta = 167.9, 158.2, 157.4, 157.2, 150.8, 145.3,$ 123.6, 119.5, 118.9, 61.5, 42.7, 40.7, 34.7, 14.0. HRMS (FAB) Calcd for C₁₄H₁₈N₆O₃ (M+H⁺) 319.1518. Found 319.1534.

Synthesis of compound 7

A solution of triphenylphosphine (66 mg, 0.22 mmol), palladium acetate (40 mg, 0.18 mmol), and triethylamine (1.05 mL) in dry DMF was stirred at 60 °C under N₂ atmosphere until the color of the solution turned red. Then, 1-bromopyrene (320 mg 1.14 mmol) dissolved in dry DMF was added to the reaction mixture, followed by Compound **6** (280 mg, 0.88 mmol) in dry DMF. The solution was refluxed at 110 °C for 3 h. The solution was filtered using Cerite to remove the insoluble solid. After the evaporation, the residue was purified by silica gel column chromatography (CHCl₃:MeOH = 20:1) to give **7** (220 mg, 48%). ¹H-NMR [DMSO-d₆, 500 MHz] δ = 11.42 (br, 1H), 8.74 (d, 1H), 8.65-8.10 (m, 10 H), 7.58 (d, 1H), 5.29 (s, 1H), 4.21 (s, 2H), 3.17 (s, 3H), 3.05 (s, 3H), 1.24 (t, 3H). ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 168.6, 158.6,

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157.9, 157.6, 151.5, 146.6, 131.5, 131.4, 130.8, 130.5, 129.2, 128.7, 128.6, 128.2, 127.9, 127.0, 126.2, 125.9, 125.7, 124.8, 124.5, 124.3, 123.0, 120.1, 117.3, 61.9, 43.4, 41.2, 35.2, 14.5. HRMS (FAB) Calcd for $C_{30}H_{26}N_6O_3$ (M+H⁺): 519.2144. Found 519.2173.

Synthesis of compound 8

Sodium hydroxide (34 mg, 0.84 mmol) in a H₂O:EtOH = 1:1 solution was added to a stirred solution of Compound **7** (220 mg, 0.42 mmol) in 1,4-dioxane:MeOH = 2:1 solution. The reaction mixture was stirred for 20 min at room temperature. After the addition of a large amount of H₂O, the pH was adjusted to 2-3 using 1 N HCl aq. The solid was collected by filtration using diethyl ether to give **8** (150 mg, 86%). ¹H-NMR [DMSO-d₆, 500 MHz] δ = 11.63 (br, 1H), 8.86 (d, 1H), 8.73-8.79 (m, 10H), 7.62 (d, 1H), 5.22 (s, 1H), 3.19 (s, 3H), 3.06 (s, 3H). ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 169.1, 158.2, 157.5, 156.5, 150.5, 146.1, 131.4, 131.0, 130.3, 129.4, 128.5, 128.4, 127.4, 126.6, 125.9, 125.6, 125.3, 124.3, 123.9, 122.4, 118.1, 66.4, 43.2, 40.9. HRMS (FAB) Calcd for C₂₈H₂₂N₆O₃ (M+H⁺): 491.1831. Found 491.1836.

Synthesis of compound 9

To a stirred solution of compound 8 (0.15 g, 0.36 mmol) and triethylamine (three drops) dissolved in DMF, 1M DMTr-Lserinol¹⁰ in DMF (0.33 mL, 0.33 mmol) was added. 4-(4,6dimethoxy-1,3,5-triazin-2-yl)-4-methylmorpholinium chloride (DMT-MM) (0.16 g, 0.50 mmol) was added to the solution at 0 °C, and stirred for 20 min at 0 °C, followed by an additional hour at room temperature. 20 mL CHCl₃ was added to the solution and stirred for 10 min. The solution was extracted with CHCl₃ and sat. NaHCO₃ aq twice. After evaporation, the residue was purified by silica gel column chromatography (CHCl₃:MeOH = 20:1, 1% triethylamine) to give **9** (190 mg, 61%). ¹H-NMR [DMSO-d_6, 500 MHz] δ = 11.38 (br, 1H), 8.71-8.08 (m, 12H), 7.43-7.13 (m, 9H), 6.75 (m, 4H), 5.06 (dd, 2H), 4.81 (t, 1H), 4.07 (m, 1H), 3.642, 3.638 (s*2, 3H), 3.54 (m, 2H), 3.03 (m, 2H), 2.98 (s, 3H), 3.82 (s, 3H). ¹³C-NMR [DMSO-d₆, 125 MHz] δ = 166.4, 157.93, 157.90, 157.5, 157.0, 151.3, 146.3, 145.0, 135.8, 135.7, 131.0, 130.8, 130.4, 130.1, 129.7, 129.6, 128.4, 128.2, 128.1, 127.73, 127.66, 127.4, 126.5, 125.7, 125.4, 125.2, 124.3, 124.1, 123.6, 122.5, 119.7, 117.2, 113.0, 85.2, 62.4, 60.8, 55.0, 54.9, 51.3, 44.3, 34.6. HRMS (FAB) Calcd for C₅₂H₄₇N₇O₆ (M+H⁺): 866.3666 Found 866.3697.

Synthesis of compound 10

Compound **9** (190 mg, 0.22 mmol) and triethylamine (0.13 mL, 1.1 mmol) were dissolved in dry dichloromethane under nitrogen. 2-cyanoethyl diisopropylchlorophosphoramidite (70 μ L, 0.33 mmol) was added dropwise to the solution at 0 °C. After stirring the mixture at room temperature for 30 min, it was subjected to silica gel column chromatography (CHCl₃:acetone = 3:1 containing 3% triethylamine). The products were then dissolved in a small amount of dry CHCl₃ and re-precipitated three times from hexane, yielding compound **10** (170 mg, 73%). Prior to synthesis of SNA on a DNA synthesizer, compound **10** was dried by co-evaporation with mixture of dry acetonitrile and dry dichloromethane. ¹H-NMR [DMSO-d₆, 500 MHz] δ =

11.39 (br, 1H), 8.72-8.10 (m, 12H), 7.42-7.14 (m, 10H), 6.76 (m, 4H), 5.08 (d, 2H), 4.19 (m, 1H), 4.07 (m, 1H), 3.70 3.55 (H) 900H), 3.42 (m, 1H), 3.10 (m, 2H), 2.99,2.98 (s*2, 3H), 2.83,2.80 (s*2, 3H), 2.61 (m, 2H), 1.02 (m, 6H), 0.93 (m, 3H). ¹³C-NMR [DMSOd₆, 125 MHz] δ = 166.4, 158.0, 157.9, 157.8, 157.5, 157.0, 151.2, 146.3, 144.9, 135.6, 135.5, 131.0, 130.8, 130.4, 130.1, 129.7, 129.6, 128.4, 128.2, 128.1, 127.7, 127.6, 127.4, 126.5, 125.7, 125.4, 125.2, 124.3, 124.0, 123.5, 122.5, 119.7, 118.8, 117.3, 113.0, 85.2, 62.0, 58.2, 58.1, 54.9, 44.2, 42.4, 42.3, 34.6, 32.1, 29.6, 24.3, 24.2, 24.1, 19.8, 19.7. ³¹P-NMR [DMSOd₆, 202 MHz] δ = 146.9, 146.6. HRMS (FAB) Calcd for C₆₁H₆₄N₉O₇P (M+H⁺) 1066.4744. Found 1066.4730.

Synthesis and purification of oligonucleotides

SNA phosphoramidite monomers involving T, G, A, and C were synthesized following reported procedures.⁹ SNA oligomers (SNA-2^{PV}G and SNA-1^{PV}G) were synthesized using an NTS M-6-MX_A12N DNA/RNA synthesizer (Nihon Techno Service Co., Ltd.) using phosphoramidite chemistry. For the coupling of all SNA monomers, the incubation time was extended to 600 s. The concentrations of the SNA phosphoramidite monomers were adjusted to 0.1 M for **10** (^{PV}G) and G, 0.75 M for T, A, and C. **10** (^{PV}G) was dissolved in a dry CH₂Cl₂:acetonitrile = 1:1 solution. The synthesized oligonucleotides were purified using Poly-Pak cartridges. The collected residue was further purified by reversed-phase HPLC (Kanto Chemical, Mightysil RP-18 GPII column). After purification, the synthesized oligonucleotides were characterized by MALDI-TOF MS. The preparation of SNA-2^{PV}A was described in a previous report.^{11a}

Conclusions

We synthesized a ^{PV}G-linked SNA for the photoresponsive system. Previous report has shown that PVA-SNA favored the formation of intrastrand photodimers upon irradiation with blue light. In contrast, PVG selectively formed interstrand photodimers at a slower reaction rate, likely due to the different orientation favouring photocycloaddition. The interstrand crosslinking and cycloreversion of SNA-2^{PV}G/RNA-2C enable the photoregulation of the dissociation and formation of duplexes upon light irradiation. The formation of by-products during irradiation of the SNA-2PVG/RNA-2C duplex may prevent the photoreaction from being reversible. In contrast, the interstrand photocycloaddition and cycloreversion of singlestranded SNA-1^{PV}G were sufficiently fast and effective compared to the SNA-2^{PV}G/RNA-2C duplex. Ogasawara et al. reported a photoregulation system using the trans-cis photoisomerization of ^{PV}G-DNA, but the photocrosslinking property of ^{PV}G was not described.¹² The rigid cyclic structure of the DNA scaffold could suppress photocycloaddition, while the flexible acyclic SNA scaffold probably enables interstrand photocycloaddition with high efficiency. As we have demonstrated, effective and reversible interstrand crosslinking between ^{PV}Gs is useful as a component of photoreactive materials such as photocaging system. In particular, ^{PV}G-SNA is a suitable candidate as a photocrosslinker for various materials that respond to visible light, instead of other

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photocrosslinkers,¹⁸ because ^{PV}G-SNA facilitates effective intermolecular photocrosslinking even in μ M concentrations due to strong stacking interactions. The powerful photocycloaddition properties of ^{PV}G-SNA promise applications in new photoresponsive nanodevices and advanced applications, such as chemical artificial intelligence.¹⁹

Author contributions

K.M. and H.A. conceived and designed the experiments. A.I. performed all experiments and F.K. instructed her organic syntheses. All authors analyzed the data, discussed the results, and co-wrote the manuscript.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the Supplementary Information.

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The data supporting this article have been included as part of the Supplementary Information.