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As featured in:



See Yusuke Takezawa, Mitsuhiro Shionoya *et al.*, *Chem. Sci.*, 2024, **15**, 2365.

Cite this: *Chem. Sci.*, 2024, 15, 2365

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# Ligase-mediated synthesis of Cu<sup>II</sup>-responsive allosteric DNAzyme with bifacial 5-carboxyuracil nucleobases†

Yusuke Takezawa,<sup>✉</sup> Hanci Zhang, Keita Mori, Lingyun Hu and Mitsuhiro Shionoya<sup>✉</sup>

A Cu<sup>II</sup>-responsive allosteric DNAzyme has been developed by introducing bifacial 5-carboxyuracil (caU) nucleobases, which form both hydrogen-bonded caU–A and metal-mediated caU–Cu<sup>II</sup>–caU base pairs. The base sequence was logically designed based on a known RNA-cleaving DNAzyme so that the caU-modified DNAzyme (caU-DNAzyme) can form a catalytically inactive structure containing three caU–A base pairs and an active form with three caU–Cu<sup>II</sup>–caU pairs. The caU-DNAzyme was synthesized by joining short caU-containing fragments with a standard DNA ligase. The activity of caU-DNAzyme was suppressed without Cu<sup>II</sup>, but enhanced 21-fold with the addition of Cu<sup>II</sup>. Furthermore, the DNAzyme activity was turned on and off during the reaction by the addition and removal of Cu<sup>II</sup> ions. Both ligase-mediated synthesis and Cu<sup>II</sup>-dependent allosteric regulation were achieved by the bifacial base pairing properties of caU. This study provides a new strategy for designing stimuli-responsive DNA molecular systems.

Received 25th September 2023

Accepted 17th January 2024

DOI: 10.1039/d3sc05042d

rsc.li/chemical-science

## Introduction

The highly sophisticated molecular recognition abilities of nucleic acids based on complementary hydrogen-bonded base pairing have led to a dramatic growth in the research field recognized as DNA nanotechnology.<sup>1</sup> Numerous DNA nano-devices, sensors, and molecular machines have been created by controlling DNA hybridization and structures in response to stimuli such as DNA/RNA binding, pH changes, and light irradiation.<sup>2</sup> Metal ions also serve as external stimuli to regulate the DNA structure and function, particularly by exploiting metal-mediated unnatural base pairing.<sup>3</sup> Metal-mediated artificial base pairs are formed between two opposing ligand-type nucleobase analogs by complexation with a bridging metal ion. Metal-mediated base pairing generally stabilizes DNA duplexes, thus controlling DNA hybridization in a metal-dependent manner.

Aiming to switch DNA functions efficiently by metal complexation, we have recently established a new concept of metal-mediated base pair switching of bifacial 5-modified pyrimidine nucleobases.<sup>4–7</sup> Bifacial bases such as 5-hydroxyuracil (U<sup>OH</sup>)<sup>4,5</sup> and 5-carboxyuracil (caU)<sup>6</sup> are designed to form metal-mediated self-base pairs (e.g., U<sup>OH</sup>–Gd<sup>III</sup>–U<sup>OH</sup>) in the

presence of certain metal ions and to form Watson–Crick-like base pairs with a natural nucleobase in DNA duplexes (e.g., U<sup>OH</sup>–A). Based on the switching between U<sup>OH</sup>–A and U<sup>OH</sup>–Gd<sup>III</sup>–U<sup>OH</sup> base pairs, Gd<sup>III</sup>-triggered DNA strand displacement reactions were demonstrated and Gd<sup>III</sup>-mediated control of DNA tweezer structures and DNAzyme functions was successfully achieved.<sup>5</sup>

In this study, a metal-responsive DNAzyme was newly developed by utilizing bifacial 5-carboxyuracil (caU) bases as metal binding sites (Fig. 1). DNAzymes are DNA molecules with catalytic activity that have been widely applied for the

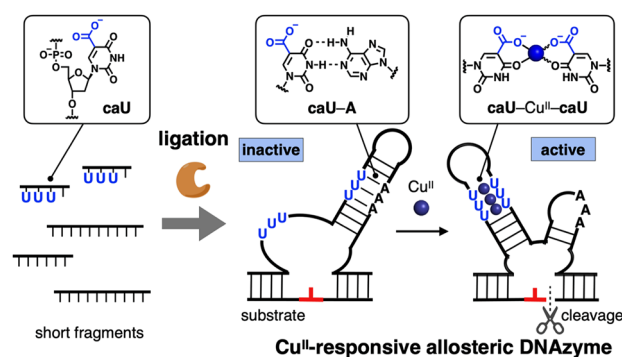


Fig. 1 Ligase-mediated synthesis of a Cu<sup>II</sup>-responsive DNAzyme containing bifacial 5-carboxyuracil (caU) bases, which form hydrogen-bonded caU–A and metal-mediated caU–Cu<sup>II</sup>–caU base pairs. U in the figure represents a caU base.

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3sc05042d>

development of DNA-based biosensors and molecular machines.<sup>8</sup> Of particular interest is the rational design of allosteric DNazymes whose activity can be controlled in response to specific stimuli. Such stimuli-responsive DNazymes are versatile components for building up deformable DNA nano-architectures as well as DNA reaction networks. Metal-responsive DNazymes have been developed previously by incorporating metal-mediated base pairs such as Cu<sup>II</sup>-mediated hydroxypyridone base pairs (H-Cu<sup>II</sup>-H).<sup>9</sup> The bifacial **caU** bases used in this study not only form hydrogen-bonded **caU**-**A** base pairs, but also Cu<sup>II</sup>-mediated **caU**-Cu<sup>II</sup>-**caU** base pairs with high metal selectivity.<sup>6</sup> The **caU**-Cu<sup>II</sup>-**caU** base pairing significantly stabilizes the DNA duplexes ( $\Delta T_m = +30.7$  °C for a duplex with three **caU**-Cu<sup>II</sup>-**caU** pairs), whereas the **caU**-**A** base pairs are destabilized by the addition of Cu<sup>II</sup> ions ( $\Delta T_m = -7.3$  °C for a duplex with three **caU**-**A** pairs by using 6 equiv. of Cu<sup>II</sup> ions). Therefore, carefully designed DNazymes containing **caU** bases were expected to exhibit good responsiveness to Cu<sup>II</sup> ions. To prepare DNzyme strands containing multiple **caU** nucleotides, enzymatic synthesis using a DNA ligase was also investigated. A standard T4 DNA ligase was utilized because it was reported to tolerate modified base pairs and backbones.<sup>9b,d,10</sup> Thus, we expected that **caU**-containing short fragments could be enzymatically ligated to give long DNA strands modified with **caU** bases.

## Results and discussion

The Cu<sup>II</sup>-responsive allosteric DNzyme was logically designed by modifying the base sequence of the known RNA-cleaving NaA43 DNzyme<sup>11</sup> (Fig. 2a) in a manner similar to the Gd<sup>III</sup>-responsive DNzyme with U<sup>OH</sup> bases.<sup>5</sup> The NaA43 DNzyme was chosen because it does not require metal cofactors that can be trapped by common chelators used to selectively remove Cu<sup>II</sup> ions for reversible regulation of DNzyme activity (*vide infra*). Since duplex stabilization is more pronounced when three or more consecutive **caU**-Cu<sup>II</sup>-**caU** pairs are used,<sup>6</sup> we decided to incorporate three **caU**-Cu<sup>II</sup>-**caU** pairs into the parent NaA43 DNzyme. Three pairs of **caU** bases were introduced into the stem region and the surrounding bases (shown in orange) were redesigned to form a different secondary structure in the absence of Cu<sup>II</sup> ions. The **caU**-modified DNzyme (**caU**-DNzyme) was expected to undergo a structural change upon addition of Cu<sup>II</sup> ions, from a catalytically inactive structure with

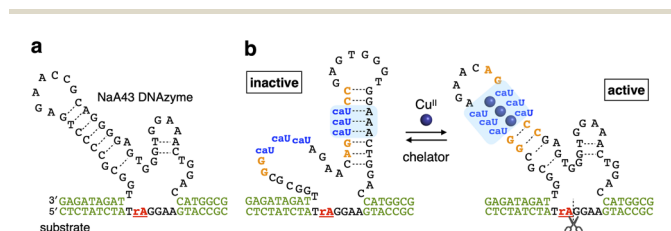


Fig. 2 Molecular design of a Cu<sup>II</sup>-responsive allosteric DNzyme with **caU** nucleobases (**caU**-DNzyme). (a) Sequence of the parent RNA-cleaving DNzyme (NaA43 DNzyme). (b) Sequence of **caU**-DNzyme. Both the active structure with **caU**-Cu<sup>II</sup>-**caU** base pairs and the inactive structure with **caU**-**A** base pairs are shown. "ra" in the substrate indicates an adenosine ribonucleotide at the cleavage site.

three **caU**-**A** base pairs to an active form with three **caU**-Cu<sup>II</sup>-**caU** base pairs (Fig. 2b). The plausible secondary structure was simulated using the NUPACK software<sup>12</sup> (Fig. S1†). The **caU** bases were replaced with natural T bases to calculate the structure in the absence of Cu<sup>II</sup>, and the potential **caU**-Cu<sup>II</sup>-**caU** pairs were changed to G-C base pairs to simulate the structure in the presence of Cu<sup>II</sup>. The results indicated that both the inactive and the active structures can be stably formed *via* the formation of **caU**-**A** and **caU**-Cu<sup>II</sup>-**caU** base pairs, respectively.

A **caU**-modified DNzyme strand (58-nt) was synthesized by ligating short DNA fragments (Fig. 3a). The **caU**-containing strands can be chemically synthesized based on the conventional phosphoramidite chemistry,<sup>6</sup> but the chemical synthesis requires an additional protecting group on the carboxylate of the **caU** bases. Although the coupling yield is sufficiently high, incomplete deprotection often makes purification of long oligonucleotides containing multiple **caU** bases very difficult. Therefore, we expected that the ligase-mediated synthesis would be a suitable strategy to synthesize **caU**-modified DNzymes. Since **caU** nucleobases can form **caU**-**A** base pairs with adenine bases on the complementary DNA, **caU**-modified oligonucleotides were expected to assemble on the splint strand and to be ligated by a standard T4 DNA ligase. A DNA tetramer 5'-**caUcaUcaUG**-3' (**1**) containing three **caU** nucleotides was used to introduce three consecutive **caU** bases into the resulting DNA strands. A natural nucleotide G was added at the 3'-terminal so that an additional G-C base pairing would facilitate the hybridization of tetramer **1** to the splint DNA **5**. The **caU**-containing fragment **1** was prepared using an automated DNA synthesizer according to the reported procedure.<sup>6</sup> Prior to the ligation reaction, fragments **1**, **2**, and **3** were treated with T4 polynucleotide kinase (T4 PNK) to introduce a phosphate group at the 5' end (step i). After all the DNA fragments were hybridized to splint **5** (step ii), a ligation reaction was performed using T4 DNA ligase (step iii). Reaction products were analyzed by

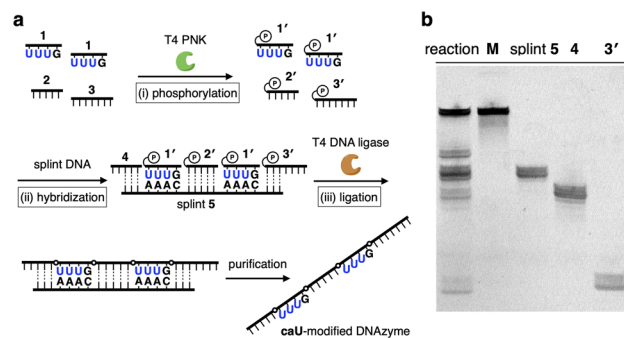


Fig. 3 Ligase-mediated synthesis of DNA strands containing **caU** nucleobases. (a) Reaction scheme. (i) A short DNA strand **1** containing three **caU** bases (5'-**caUcaUcaUG**-3') was phosphorylated by T4 polynucleotide kinase (T4 PNK). (ii) DNA fragments were hybridized with a splint DNA strand **5**. (iii) The DNA fragments were ligated by T4 DNA ligase. [1'] = 30 μM, [2'] = [3'] = 12 μM, [4] = [5] = 10 μM, [ligase] = 2 U per μL in quick ligation buffer, 16 °C, 18 h. U in the figure represents a **caU** base. (b) Denaturing PAGE analysis of the reaction products. The bands were detected after SYBR gold staining. M: chemically synthesized T-DNzyme, which has T bases instead of **caU** bases.



denaturing polyacrylamide gel electrophoresis (PAGE) (Fig. 3b). After incubation at 16 °C for 18 h, fragment **4** was efficiently consumed and a low mobility band appeared. The mobility of the new band was nearly identical to that of a chemically synthesized T-DNAzyme strand (58-nt) in which the **caU** nucleotides of the **caU**-DNAzyme were replaced with natural thymidines (T). Comparing the band intensities, the reaction yield reached 70% or more, confirming that the ligase reaction was proceeding well. The formation of the desired **caU**-DNAzyme strand was confirmed by MALDI mass spectrometry after isolation ( $[M-2H]^{2-}$ : calcd 9150.31, found 9150.67, Fig. S2†). It was demonstrated that T4 PNK and T4 DNA ligase successfully phosphorylated and ligated the short strand **1** (5'-**caUcaUcaUG**-3') despite the presence of a modified **caU** base at the 5' end. Since the desired product can be easily isolated by denaturing PAGE, the ligase-mediated synthesis was shown to be a powerful alternative method to incorporate multiple **caU** bases into functional DNA sequences.

Using the resulting **caU**-modified DNAzyme strand, we examined whether the catalytic activity of **caU**-DNAzyme can be regulated in response to the addition of  $\text{Cu}^{\text{II}}$  ions. DNAzyme-catalyzed RNA-cleaving reactions were performed using 10 equiv. of substrates labeled with a fluorescent dye (FAM). Substrate cleavage was quantitatively evaluated by denaturing PAGE analysis (Fig. S3†). Fig. 4a compares the RNA cleavage reaction catalyzed by **caU**-DNAzyme in the presence of varying concentrations of  $\text{Cu}^{\text{II}}$  ions. In the absence of  $\text{Cu}^{\text{II}}$  ions, the RNA-cleaving activity of the **caU**-DNAzyme was greatly suppressed. The DNAzyme activity was found to be enhanced by the

addition of  $\text{Cu}^{\text{II}}$  ions; the highest activity was observed when 9 equiv. of  $\text{Cu}^{\text{II}}$  ions were added. Time-course analysis further confirmed the  $\text{Cu}^{\text{II}}$ -dependent activation of **caU**-DNAzyme (Fig. 4b and S4†). Under the same conditions, the catalytic activity of the unmodified NaA43 DNAzyme was reduced by adding  $\text{Cu}^{\text{II}}$  ions (Fig. S5†). A control T-DNAzyme containing natural T bases in place of **caU** showed no RNA-cleaving activity both in the absence and presence of  $\text{Cu}^{\text{II}}$  ions. These results clearly demonstrate that the addition of  $\text{Cu}^{\text{II}}$  ions enhances the catalytic activity of the **caU**-modified DNAzyme. In the presence of  $\text{Cu}^{\text{II}}$  ions, the **caU**-DNAzyme cleaved approximately 70% of the substrate (*i.e.*, 7 equiv.) in 20 h, confirming that the modified DNAzyme maintains multiple turnover ability.

It is most likely that the activity of **caU**-DNAzyme was switched based on the changes in the base-pairing partners of the **caU** bases. This was supported by a model experiment using a FAM-labeled strand containing three **caU** bases in the middle (**1U**) (Fig. 4c and S6†). The DNA **1U** was annealed with two complementary strands **2U** and **2A** containing three **caU** or A bases, respectively, and the hybridization products were analyzed by native PAGE (Fig. 4d). Under  $\text{Cu}^{\text{II}}$ -free conditions, only the duplex **1U**·**2A** with **caU**-A pairs was formed. In the presence of  $\text{Cu}^{\text{II}}$  ions, the duplex **1U**·**2U** with **caU**- $\text{Cu}^{\text{II}}$ -**caU** pairs was formed (up to about 60%). These results clearly show the  $\text{Cu}^{\text{II}}$ -mediated change in the hybridization partners, which is the driving force behind the allosteric regulation of the **caU**-DNAzyme.

The catalytically active form contains three **caU**- $\text{Cu}^{\text{II}}$ -**caU** base pairs, but the maximum activity was observed in the presence of 9 equiv. of  $\text{Cu}^{\text{II}}$  ions (Fig. 4a). This inconsistency can be explained by the stability of the **caU**-A base pairs in addition to the **caU**- $\text{Cu}^{\text{II}}$ -**caU** pairs.<sup>6</sup> Melting experiments (Fig. S7†) showed that the model 15-bp duplex **1U**·**2U**, containing three **caU**-**caU** pairs, exhibited the highest melting temperature ( $T_m$ ) in the presence of 3 equiv. of  $\text{Cu}^{\text{II}}$  ions, due to the quantitative formation of **caU**- $\text{Cu}^{\text{II}}$ -**caU** pairs. On the other hand, the stability of duplex **1U**·**2A**' with three **caU**-A pairs decreases with increasing amounts of  $\text{Cu}^{\text{II}}$  ions, possibly due to the binding of  $\text{Cu}^{\text{II}}$  ions to the **caU** bases.<sup>6</sup> The difference in the  $T_m$  values of duplexes **1U**'·**2U** and **1U**'·**2A**' was maximal when more than 3 equiv. of  $\text{Cu}^{\text{II}}$  were added. In fact, the hybridization experiments (Fig. 4d) showed that an excess of  $\text{Cu}^{\text{II}}$  ions are required to change the hybridization partners of the **caU** bases. Therefore, the requirement for an excess of  $\text{Cu}^{\text{II}}$  ions to activate the **caU**-DNAzyme suggests that the DNAzyme functions through both  $\text{Cu}^{\text{II}}$ -mediated destabilization of the **caU**-A pairs (inactive state) and  $\text{Cu}^{\text{II}}$ -mediated **caU**- $\text{Cu}^{\text{II}}$ -**caU** base pair formation (active state) exactly as designed.

The apparent first-order rate constants ( $k_{\text{obs}}$ ) for the DNAzyme reactions were estimated from the initial rates (Table 1). The catalytic activity of **caU**-DNAzyme was found to increase by approximately 21-fold with the addition of 9 equiv. of  $\text{Cu}^{\text{II}}$  ions. The maximum activity of the **caU**-DNAzyme ( $k_{\text{obs}} = 4.2 \times 10^{-2} \text{ h}^{-1}$ ) was lower than that of the unmodified NaA43 DNAzyme ( $k_{\text{obs}} = 4.7 \times 10^{-1} \text{ h}^{-1}$ ). This may be due not only to the incomplete transformation into the active state, but also to the structural distortion caused by the **caU**- $\text{Cu}^{\text{II}}$ -**caU** base pairs.<sup>6</sup> As

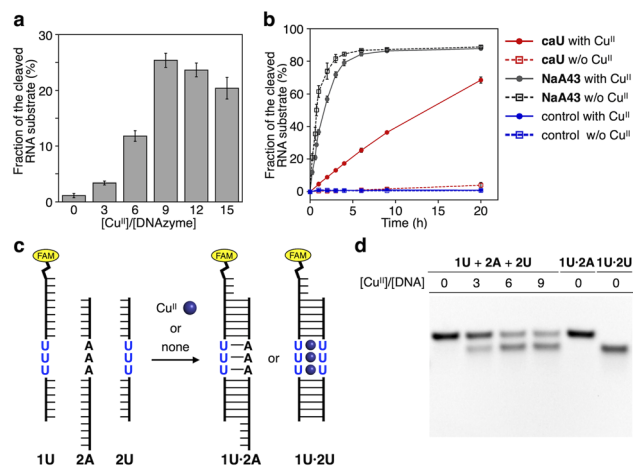


Fig. 4 (a) RNA-cleaving activity of **caU**-DNAzyme in the presence of varying concentrations of  $\text{Cu}^{\text{II}}$  ions. The fractions of the cleaved substrate after a 6 h reaction are shown (see also Fig. S4†). (b) RNA-cleaving activity of **caU**-DNAzyme and the parent NaA43 DNAzyme in the absence and the presence of 9 equiv. of  $\text{Cu}^{\text{II}}$ . T-DNAzyme, in which all **caU** bases are replaced with natural T bases, was used as a control. [DNAzyme] = 1.0  $\mu\text{M}$ , [substrate] = 10  $\mu\text{M}$ ,  $[\text{CuSO}_4] = 0, 9.0 \mu\text{M}$  in 10 mM HEPES (pH 7.0), 100 mM NaCl, 25 °C. (c)  $\text{Cu}^{\text{II}}$ -mediated change in the hybridization partners of the **caU**-containing strand (**1U**). (d) Native PAGE analysis of the hybridization product in the presence of varying amounts of  $\text{Cu}^{\text{II}}$  ions. [DNA] = 2  $\mu\text{M}$  each in 10 mM HEPES (pH 7.0) and 100 mM NaCl. The samples were annealed prior to the analysis. FAM detection.



**Table 1** Apparent first-order rate constants ( $k_{\text{obs}}$ ) for the DNAzyme-catalyzed RNA cleavage reactions

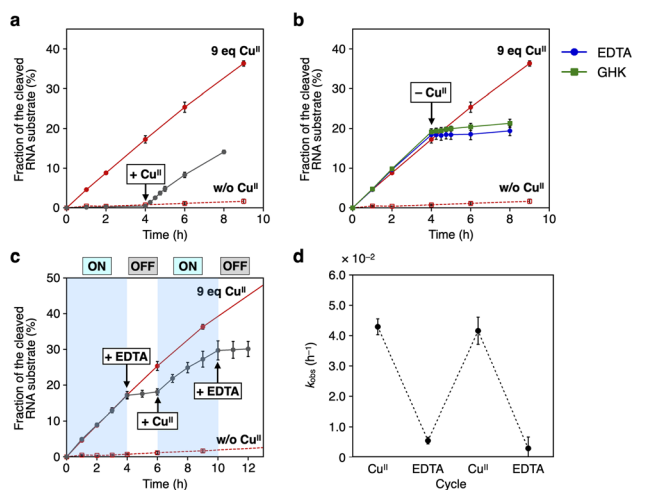
DNAzyme	$k_{\text{obs}}/\text{h}^{-1}$		Ratio
	$\text{Cu}^{\text{II}+}$	$\text{Cu}^{\text{II}-}$	
caU-DNAzyme	$4.3 \times 10^{-2\text{b}}$	$2.0 \times 10^{-3}$	21
NaA43 DNAzyme	$4.7 \times 10^{-1\text{b}}$	$8.5 \times 10^{-1}$	0.56
T-DNAzyme	$<1.0 \times 10^{-3\text{b}}$	$<1.0 \times 10^{-3}$	—
H-modified DNAzyme <sup>a</sup>	$2.8 \times 10^{-1\text{c}}$	$4.7 \times 10^{-2}$	5.9

<sup>a</sup> A NaA43 DNAzyme modified with a pair of hydroxypyridone (H) nucleobases.<sup>9b</sup> <sup>b</sup> In the presence of 9 equiv. of  $\text{Cu}^{\text{II}}$  ions. <sup>c</sup> In the presence of 1 equiv. of  $\text{Cu}^{\text{II}}$  ions.

indicated by circular dichroism (CD) analysis in the previous study,<sup>6</sup> caU– $\text{Cu}^{\text{II}}$ –caU base pairing can unwind the stem duplex to some extent. Introducing the caU base at a position more distant from the catalytic core would reduce the negative effect on the DNAzyme activity. It is noteworthy that the  $\text{Cu}^{\text{II}}$ -mediated activation of caU-DNAzyme (21-fold) was much more efficient than that of a  $\text{Cu}^{\text{II}}$ -responsive H-modified DNAzyme (5.9-fold), which was developed by incorporating an H– $\text{Cu}^{\text{II}}$ –H base pair into the same NaA43 DNAzyme. The results clearly show that the bifacial caU nucleobases, which form both hydrogen-bonded caU–A and metal-mediated caU– $\text{Cu}^{\text{II}}$ –caU base pairs, are useful for metal-responsive switching of DNA functions.

We further carried out the DNAzyme reactions in the presence of  $\text{Hg}^{\text{II}}$  ions that can mediate caU– $\text{Hg}^{\text{II}}$ –caU base pairing<sup>6</sup> (Fig. S8†). In contrast to  $\text{Cu}^{\text{II}}$  ions, the addition of  $\text{Hg}^{\text{II}}$  ions did not activate caU-DNAzyme at all. Note that the activity of the unmodified NaA43 DNAzyme was significantly reduced by the addition of  $\text{Hg}^{\text{II}}$  ions. This is probably due to the undesired binding of  $\text{Hg}^{\text{II}}$  ions to the natural T bases. These results clearly demonstrate the advantage of using metal ions that do not interact strongly with natural bases (e.g.,  $\text{Cu}^{\text{II}}$ ), especially with long oligonucleotides such as DNAzymes.

The RNA-cleaving activity of caU-DNAzyme was reversibly regulated in response to  $\text{Cu}^{\text{II}}$  ions during the reaction (Fig. 5). The reaction was initiated in the absence of  $\text{Cu}^{\text{II}}$  ions, and after 4 h,  $\text{Cu}^{\text{II}}$  ions (9 equiv.) were added. The reaction rate was immediately increased to  $k_{\text{obs}} = 3.9 \times 10^{-2} \text{ h}^{-1}$ , which is comparable to the rate observed when the reaction was initiated with  $\text{Cu}^{\text{II}}$  ions (Fig. 5a). In a similar manner, removal of  $\text{Cu}^{\text{II}}$  ions was shown to inactivate caU-DNAzyme. Addition of the chelating agent EDTA or  $\text{Cu}^{\text{II}}$ -binding peptide (GHK)<sup>13</sup> in equimolar amounts with  $\text{Cu}^{\text{II}}$  ions (9 equiv.) immediately slowed the reaction ( $k_{\text{obs}} = 2.5 \times 10^{-3} \text{ h}^{-1}$  and  $5.3 \times 10^{-3} \text{ h}^{-1}$ , respectively) (Fig. 5b). The addition of sodium ascorbate, which can reduce  $\text{Cu}^{\text{II}}$  to  $\text{Cu}^{\text{I}}$ , also decreased the activity of caU-DNAzyme to  $k_{\text{obs}} = 3.6 \times 10^{-3} \text{ h}^{-1}$  (Fig. S9†). These results demonstrate that the activity of caU-DNAzyme can be rapidly switched by the addition, removal, and reduction of  $\text{Cu}^{\text{II}}$  ions under isothermal conditions. Alternate addition of  $\text{Cu}^{\text{II}}$  ions (9 equiv.) and EDTA (9 equiv.) cycled the on–off switching of caU-DNAzyme (Fig. 5c). The  $k_{\text{obs}}$  values in each step demonstrate a clear switching in the caU-DNAzyme activity in response to  $\text{Cu}^{\text{II}}$  (Fig. 5d).



**Fig. 5**  $\text{Cu}^{\text{II}}$ -dependent regulation of the RNA-cleaving activity of caU-DNAzyme. (a) Activation of caU-DNAzyme by the addition of  $\text{Cu}^{\text{II}}$  ions (9 equiv.). (b) Deactivation of caU-DNAzyme by the removal of  $\text{Cu}^{\text{II}}$  ions with a chelating agent EDTA or a  $\text{Cu}^{\text{II}}$ -binding tripeptide (GHK) (9 equiv.). (c) Iterative switching of the DNAzyme activity.  $\text{Cu}^{\text{II}}$  (9 equiv.) and EDTA (9 equiv.) were alternately added. [DNAzyme] = 1.0  $\mu\text{M}$ , [substrate] = 10  $\mu\text{M}$ , 25 °C. The activities of caU-DNAzyme in the absence (red dotted lines) and presence of  $\text{Cu}^{\text{II}}$  ions (red solid lines) are also shown. (d) Apparent first-order rate constant ( $k_{\text{obs}}$ ) for each step.  $N = 3$ . Error bars indicate standard errors.

## Conclusions

In summary, a  $\text{Cu}^{\text{II}}$ -responsive allosteric DNAzyme was developed by introducing 5-carboxyuracil (caU) nucleobases into a known DNAzyme sequence. The caU-modified DNAzyme (caU-DNAzyme) was enzymatically synthesized by joining short caU-containing fragments with a standard T4 DNA ligase. The ligase-mediated synthesis was possible because the caU base was structurally similar to the natural T base and could form a Watson–Crick-like base pair with the A base on the splint DNA. The base sequence of caU-DNAzyme is logically designed to form both the catalytically inactive structure by caU–A base pairing and the active form by metal-mediated caU– $\text{Cu}^{\text{II}}$ –caU base pairing. The activity of caU-DNAzyme was enhanced 21-fold by the addition of  $\text{Cu}^{\text{II}}$  ions and could be turned on and off during the reaction by the addition and removal (or reduction) of  $\text{Cu}^{\text{II}}$  ions. These results demonstrate that the caU-modified DNAzyme was allosterically regulated through metal-mediated base-pair switching between caU–A and caU– $\text{Cu}^{\text{II}}$ –caU. The use of  $\text{Cu}^{\text{II}}$  is essential to induce base-pair switching of the caU base. The caU base can form other types of metal-mediated base pairs such as caU– $\text{Hg}^{\text{II}}$ –T, caU– $\text{Ag}^{\text{I}}$ –C, and caU– $\text{Cu}^{\text{II}}$ –G.<sup>6</sup> Therefore, caU-modified DNAs are expected to be further applied in constructing more complex DNA systems responsive to multiple metal ions.

This study confirms that metal-responsive DNA systems can be logically designed based on metal-mediated base-pair switching of bifacial caU nucleobases. The strategic design of caU-modified DNAzymes is expected to be applied to other types of DNAzymes<sup>14</sup> and other functional DNAs as well. Ligase-



mediated synthesis provides a simple way to incorporate **caU** bases into longer DNA sequences and is advantageous for sequence screening. Thus, it is suggested that the incorporation of bifacial **caU** bases is a powerful strategy for creating a variety of Cu<sup>II</sup>-responsive DNA molecular systems. The range of metal ions used could be expanded by developing other types of bifacial nucleobases with a different metal coordinating functionality at the 5-position of pyrimidine bases. In fact, Gd<sup>III</sup>-responsive DNA systems have been developed using cognate U<sup>OH</sup> nucleobases.<sup>5</sup> The bifacial 5-modified pyrimidine bases are expected to be introduced into DNA *via* the ligase-mediated synthesis, similar to the case with **caU**. Therefore, metal-mediated base-pair switching of bifacial nucleobases and the ligase-mediated synthesis have the potential to be versatile tools for building DNA-based stimuli-responsive systems such as biosensors, molecular machines, and computing devices. Further applications of bifacial **caU** nucleobases to metal-triggered operation of DNA nanoarchitectures and DNA logic circuits are currently under investigation.

## Data availability

All the data supporting this study are included in the main text and the ESI.†

## Author contributions

Y. T. and M. S. conceived and directed the study. H. Z. and K. M. performed the experiments and analyzed the data with the aid of Y. T. and L. H. All the authors prepared the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by JSPS KAKENHI Grant Numbers JP18H02081 and JP21H02055 to Y. T., JP21H05022 to M. S., MEXT KAKENHI Grant Numbers JP21H00384 (Molecular Engine), JP21H05866, and JP23H04399 (Molecular Cybernetics) to Y. T., and JP16H06509 (Coordination Asymmetry) to M. S., and Grant-in-Aid for JSPS Fellows to K. M. (JP21J11325) and to L. H. (JP21J11332), Japan. We would like to thank Ms Yoko Shimada for her cooperation in the experiment.

## Notes and references

- (a) N. C. Seeman and H. F. Sleiman, *Nat. Rev. Mater.*, 2018, **3**, 17068; (b) M. Madsen and K. V. Gothelf, *Chem. Rev.*, 2019, **119**, 6384–6458; (c) E. Del Grosso, E. Franco, L. J. Prins and F. Ricci, *Nat. Chem.*, 2022, **14**, 600–613.
- (a) H. Ramezani and H. Dietz, *Nat. Rev. Genet.*, 2020, **21**, 5–26; (b) S. Lu, J. Shen, C. Fan, Q. Li and X. Yang, *Adv. Sci.*, 2021, **8**, 2100328; (c) F. Wang, X. Liu and I. Willner, *Angew. Chem., Int. Ed.*, 2015, **54**, 1098–1129; (d) S. Murata, T. Toyota, S. M. Nomura, T. Nakakuki and A. Kuzuya, *Adv. Funct. Mater.*, 2022, **32**, 2201866.
- (a) Y. Takezawa, J. Müller and M. Shionoya, *Chem. Lett.*, 2017, **46**, 622–633; (b) Y. Takezawa and M. Shionoya, *Acc. Chem. Res.*, 2012, **45**, 2066–2076; (c) S. Naskar, R. Guha and J. Müller, *Angew. Chem., Int. Ed.*, 2020, **59**, 1397–1406; (d) Y. Tanaka, J. Kondo, V. Sychrovský, J. Šebera, T. Dairaku, H. Saneyoshi, H. Urata, H. Torigoe and A. Ono, *Chem. Commun.*, 2015, **51**, 17343–17360; (e) Y. Takezawa and M. Shionoya, in *Modern Avenues in Metal-Nucleic Acid Chemistry (Metal Ions In Life Science)*, vol. 25, ed. J. Müller and B. Lippert, CRC Press, Boca Raton, 2023, pp. 257–289; (f) Y. Takezawa and M. Shionoya, in *Handbook of Chemical Biology of Nucleic Acids*, ed. N. Sugimoto, Springer, Singapore, 2023, pp. 2645–2683.
- (a) Y. Takezawa, K. Nishiyama, T. Mashima, M. Katahira and M. Shionoya, *Chem.–Eur. J.*, 2015, **21**, 14713–14716; (b) K. Nishiyama, Y. Takezawa and M. Shionoya, *Inorg. Chim. Acta*, 2016, **452**, 176–180; (c) K. Nishiyama, K. Mori, Y. Takezawa and M. Shionoya, *Chem. Commun.*, 2021, **57**, 2487–2490.
- Y. Takezawa, K. Mori, W.-E. Huang, K. Nishiyama, T. Xing, T. Nakama and M. Shionoya, *Nat. Commun.*, 2023, **14**, 4759.
- Y. Takezawa, A. Suzuki, M. Nakaya, K. Nishiyama and M. Shionoya, *J. Am. Chem. Soc.*, 2020, **142**, 21640–21644.
- K. Mori, Y. Takezawa and M. Shionoya, *Chem. Sci.*, 2023, **14**, 1082–1088.
- (a) S. K. Silverman, *Trends Biochem. Sci.*, 2016, **41**, 595–609; (b) M. Liu, D. Chan and Y. Li, *Acc. Chem. Res.*, 2017, **50**, 2273–2283; (c) E. M. McConnell, I. Cozma, Q. Mou, J. D. Brennan, Y. Lu and Y. Li, *Chem. Soc. Rev.*, 2021, **50**, 8954–8994; (d) Z. Huang, X. Wang, Z. Wu and J.-H. Jiang, *Chem.–Asian J.*, 2022, **17**, e202101414.
- (a) Y. Takezawa, T. Nakama and M. Shionoya, *J. Am. Chem. Soc.*, 2019, **141**, 19342–19350; (b) T. Nakama, Y. Takezawa, D. Sasaki and M. Shionoya, *J. Am. Chem. Soc.*, 2020, **142**, 10153–10162; (c) Y. Takezawa, L. Hu, T. Nakama and M. Shionoya, *Angew. Chem., Int. Ed.*, 2020, **59**, 21488–21492; (d) T. Nakama, Y. Takezawa and M. Shionoya, *Chem. Commun.*, 2021, **57**, 1392–1395; (e) S. C. Rajasree, Y. Takezawa and M. Shionoya, *Chem. Commun.*, 2023, **59**, 1006–1009; (f) Y. Takezawa, L. Hu, T. Nakama and M. Shionoya, *Chem. Commun.*, 2024, **60**, 288–291.
- (a) R. Hili, J. Niu and D. R. Liu, *J. Am. Chem. Soc.*, 2013, **135**, 98–101; (b) D. Kong, W. Yeung and R. Hili, *J. Am. Chem. Soc.*, 2017, **139**, 13977–13980; (c) Y. Lei, J. Washington and R. Hili, *Org. Biomol. Chem.*, 2019, **17**, 1962–1965; (d) D. Kestemont, M. Renders, P. Leonczak, M. Abramov, G. Schepers, V. B. Pinheiro, J. Rozenski and P. Herdewijn, *Chem. Commun.*, 2018, **54**, 6408–6411; (e) J. Riedl, Y. Ding, A. M. Fleming and C. J. Burrows, *Nat. Commun.*, 2015, **6**, 8807; (f) N. Sabat, A. Stämpfli, S. Hanlon, S. Bisagni, F. Sladojevich, K. Püntener and M. Hollenstein, *ChemRxiv*, 2023, DOI: [10.26434/chemrxiv-2023-vwwt5](https://doi.org/10.26434/chemrxiv-2023-vwwt5).
- S.-F. Torabi, P. Wu, C. E. McGhee, L. Chen, K. Hwang, N. Zheng, J. Cheng and Y. Lu, *Proc. Natl. Acad. Sci. U. S. A.*, 2015, **112**, 5903–5908.



- 12 J. N. Zadeh, C. D. Steenberg, J. S. Bois, B. R. Wolfe, M. B. Pierce, A. R. Khan, R. M. Dirks and N. A. Pierce, *J. Comput. Chem.*, 2011, **32**, 170–173.
- 13 L. Pickart, J. H. Freedman, W. J. Loker, J. Peisach, C. M. Perkins, R. E. Stenkamp and B. Weinstein, *Nature*, 1980, **288**, 715–717.
- 14 (a) R. R. Breaker and G. F. Joyce, *Chem. Biol.*, 1995, **2**, 655–660; (b) J. W. Liu, A. K. Brown, X. L. Meng, D. M. Cropek, J. D. Istok, D. B. Watson and Y. Lu, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **104**, 2056–2061.

