



Showcasing research from Professor Mitsuhiro Shionoya's laboratory, Department of Chemistry, Graduate School of Science, The University of Tokyo, Tokyo, Japan.

Ligase-mediated synthesis of  $\text{Cu}^{\text{II}}$ -responsive allosteric DNAzyme with bifacial 5-carboxyuracil nucleobases

A  $\text{Cu}^{\text{II}}$ -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 so that the **caU**-modified DNAzyme can form a catalytically inactive structure containing **caU-A** base pairs and an active form with **caU-Cu<sup>II</sup>-caU** pairs. The **caU**-modified DNAzyme was synthesized by joining short **caU**-containing fragments with a standard DNA ligase. Both ligase-mediated synthesis and  $\text{Cu}^{\text{II}}$ -dependent allosteric regulation were achieved by the bifacial base pairing properties of **caU** nucleobases.

As featured in:



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