## **Supporting Information**

## A Label-free and enzyme-free system for operating various logic circuits using poly(thymine)-templated CuNPs and SYBR Green I as signal transducers

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Strands	Sequences ( 5'- 3')
P-DNA	CCAGA TATTC ACCGG
D <sub>0</sub>	ΑΑΑΑΑ ΑΑΑΑΑ ΑΑΑΑΑ ΑΑΑΑΑ ΑΑΑΑΑ
D1	TTTTT TTTTT TTTTT TTTTT TTTTT
D <sub>2</sub>	CCGGT GAATA TCTGG
D <sub>3</sub>	CCGGT GAATA TCTGG TTTTT TTTTT TTTTT TTTTT TTTTT TTTTT

Table S1. DNA sequences used in the designed logic operations.

To find the available DNA sequences for the developed higher-order logic gates, the designed DNA sequences were firstly mimicked on the website http://mfold.rna.albany.edu/?q=DINAMelt/Twostate-melting and then modified according to the experimental results. The above procedures were repeated until the available DNA sequences were obtained.

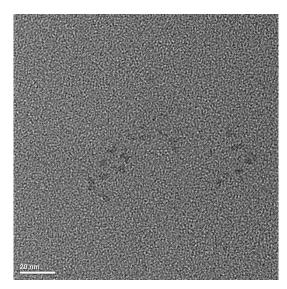


Fig. S1 Representative TEM image of as-synthesized CuNPs using polyT (T30) ss-DNA as template.

The as-synthesized CuNPs was characterized with Transmission electron microscopy (TEM) as shown in Fig. S1. The size of as-synthesized CuNPs was about 4 nm, which is consistent with previous report.<sup>1</sup>

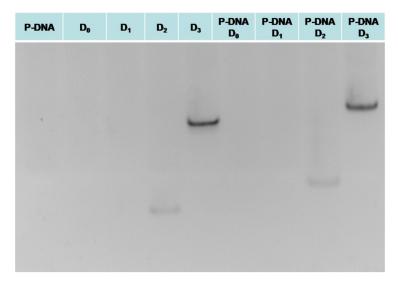


Fig. S2 PAGE analysis of DNA interactions for the 4-to-2 encoder

The first five belts from left indicate the ss-DNAs of P-DNA,  $D_0$ ,  $D_1$ ,  $D_2$  and  $D_3$  in sequence. No obvious belts can be found from P-DNA. It maybe attributes to the short length of P-DNA. The same phenomenon appears to  $D_0$  and  $D_1$  since these two poly-DNA strands cannot form any secondary structures, resulting in low binding affinity to the ds-DNA dyes used for the PAGE analysis. In the coexistence of P-DNA/ $D_0$  or the coexistence of P-DNA/ $D_1$ , no belt appears at new position, suggesting no hybridization reaction happens. Belts appear at new positions in the presence of P-DNA/ $D_2$  and P-DNA/ $D_3$ , indicating the formation of the corresponding duplex.

## References

1. Z. Qing, X. He, D. He, K. Wang, F. Xu, T. Qing and X. Yang, *Angew. Chem.Int. Ed.*, 2013, **52**, 9719-9722.