

## Electronic Supplementary Information

# A Label-Free Aptasensor for Highly Efficient ATP Detection by Using Exonuclease I and Oligonucleotide-templated Fluorescent Copper Nanoparticles

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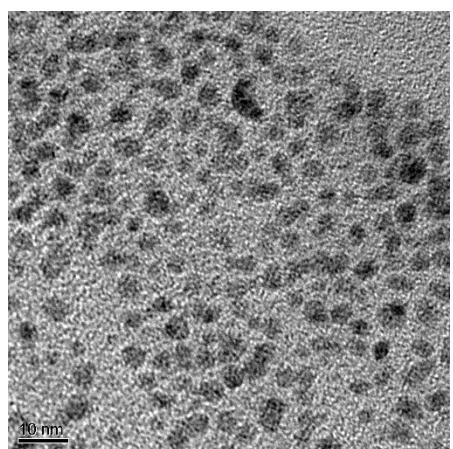
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**Fig. S1.** Typical TEM image of oligonucleotide-templated copper nanoparticles.



**Table S1** Comparision of different methods for ATP determination.

<b>Method</b>	<b>Range</b>	<b>LOD</b>	<b>Reference</b>
MB <sup>1</sup> -based assay	0.8 – 80 μM	0.5 μM	[1]
GO <sup>2</sup> -based molecular beacon assay	5 – 2500 μM	2 μM	[2]
AuNPs <sup>3</sup> -based aptamer	4.4 – 132.7 μM	0.6 μM	[3]
UCNPs <sup>4</sup> -based biosensor	0.1 – 0.75 mM	20 μM	[4]
Sandwich-type FRET <sup>5</sup> assay	2 – 16 μM	1.70 μM	[5]
]Aptamer–target recognition based aptasensor	62.5 – 2500 μM	1.45 μM	[6]
AIE <sup>6</sup> -active probe	0 – 1 mM	24 μM	[7]
Microfluidic paper analysis	0.5 – 10 μM	1 μM	[8]
GOx signaling trigger	10 – 100 μM	10 μM	[9]
Fe <sup>3+</sup> - fluorescence carbon dots	0.5 – 50 μM	0.48 μM	[10]
Zn <sup>2+</sup> -Cysteine capped CdTe QDs <sup>7</sup>	5 – 50 μM	2.07 μM	[11]
Aptamer fluorescence anisotropy sensors	1 – 200 μM	1 μM	[12]
Oligonucleotide-based CuNPs system	1 – 80 μM	0.5 μM	This work

<sup>1</sup>MB, Molecular Beacon

<sup>2</sup>GO, Graphene Oxide

<sup>3</sup>AuNPs, Gold nanoparticles

<sup>4</sup>UCNPs, Upconversion nanpparticles

<sup>5</sup>FRET, Fluorescence resonance energy transfer

<sup>6</sup>AIE, Aggregation-induced emission

<sup>7</sup>QDs, Quantum dots

## Reference

- [1] S.J. Zhen, L.Q. Chen, S.J. Xiao, Y.F. Li, P.P. Hu, L. Zhan, L. Peng, E.Q. Song, C.Z. Huang, *Anal. Chem.* 82 (2010) 8432-8437.
- [2] Y. He, Z.-G. Wang, H.-W. Tang, D.-W. Pang, *Biosens. Bioelectron.* 29 (2011) 76-81.
- [3] J. Wang, L. Wang, X. Liu, Z. Liang, S. Song, W. Li, G. Li, C. Fan, *Adv. Mater.* 19 (2007) 3943-3946.
- [4] K. Song, X. Kong, X. Liu, Y. Zhang, Q. Zeng, L. Tu, Z. Shi, H. Zhang, *Chem. Commun.* 48 (2012) 1156-1158.
- [5] X. He, Z. Li, X. Jia, K. Wang, J. Yin, *Talanta*, 111 (2013) 105-110.
- [6] Y. Tan, X. Zhang, Y. Xie, R. Zhao, C. Tan, Y. Jiang, *Analyst*, 137 (2012) 2309-2312.
- [7] K. Ma, H. Wang, H. Li, S. Wang, X. Li, B. Xu, W. Tian, *Sens. Actuators. B. Chem.* 230 (2016) 556-558.
- [8] S.-Q. Jin, S.-M. Guo, P. Zuo, B.-C. Ye, *Biosens. Bioelectron.* 63 (2015) 379-383.
- [9] S. Sitaula, S.D. Branch, M.F. Ali, *Chem. Commun.* 48 (2012) 9284-9286.
- [10] Z. Zhan, J. Cai, Q. Wang, Y. Su, L. Zhang, Y. Lv, *Luminescence*, 31 (2016) 626-632.
- [11] F. Shi, Y. Li, Z. Lin, D. Ma, X. Su, *Sens. Actuators. B. Chem.* 220 (2015) 433-440.
- [12] Q. Zhao, Q. Lv, H. Wang, *Biosens. Bioelectron.* 70 (2015) 188-193.