

Electronic Supplementary Material (ESI) for Analyst.
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Supporting Information for

**A simple molecular beacon with duplex-specific
nuclease amplification for detection of microRNA**

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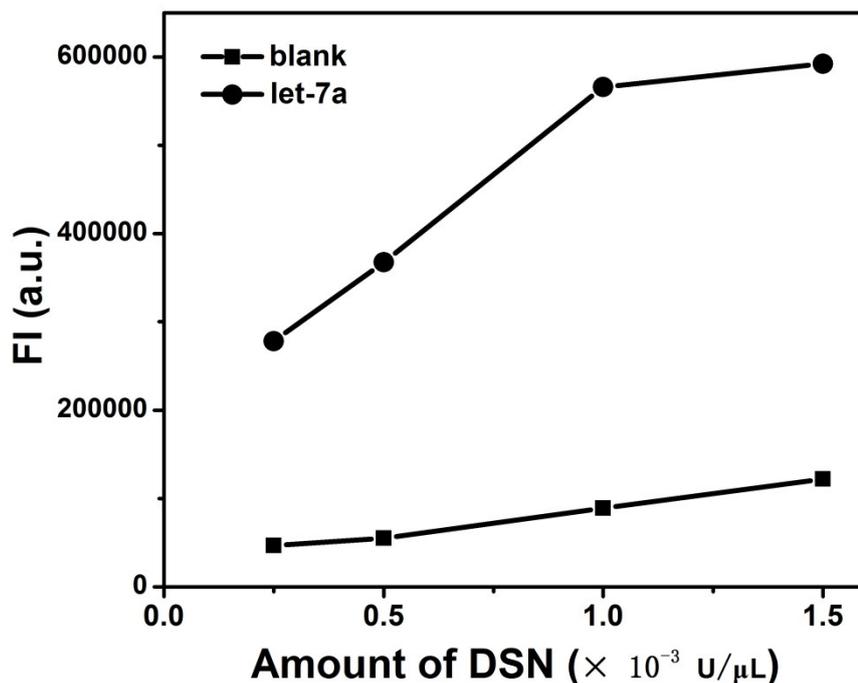


Fig. S1 The effect of DSN amount. The concentration of let-7a in the blank and the sample was 0 pM, 250 pM respectively. The reaction volume was 10 μ L and reaction products were diluted by 20 times with TE buffer solution (10 mM Tris-HCl (pH 8.0) 1 mM EDTA) before fluorescence measurement.

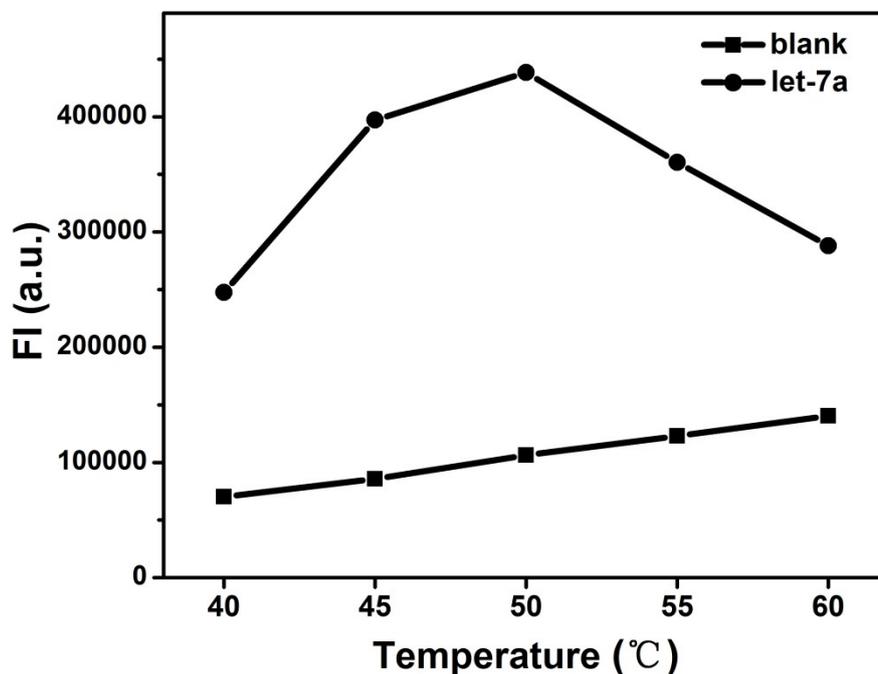


Fig. S2 The effect of the reaction temperature. The concentration of let-7a in the blank and the sample was 0 pM, 250 pM respectively. The reaction volume was 10 μ L and reaction products were diluted by 20 times with TE buffer solution before fluorescence measurement.

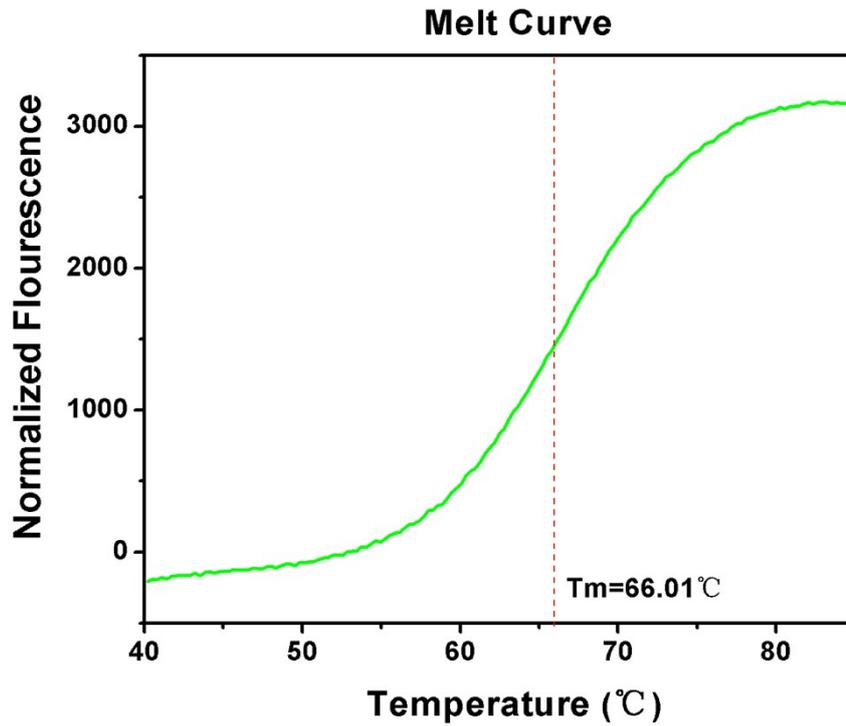


Fig. S3 Melting temperature (T_m) measurement of MB probe. T_m measurement was performed in the StepOne Real Time PCR System. The concentration of probe mb-a1 was 200 nM in 10 μ L reaction solution.

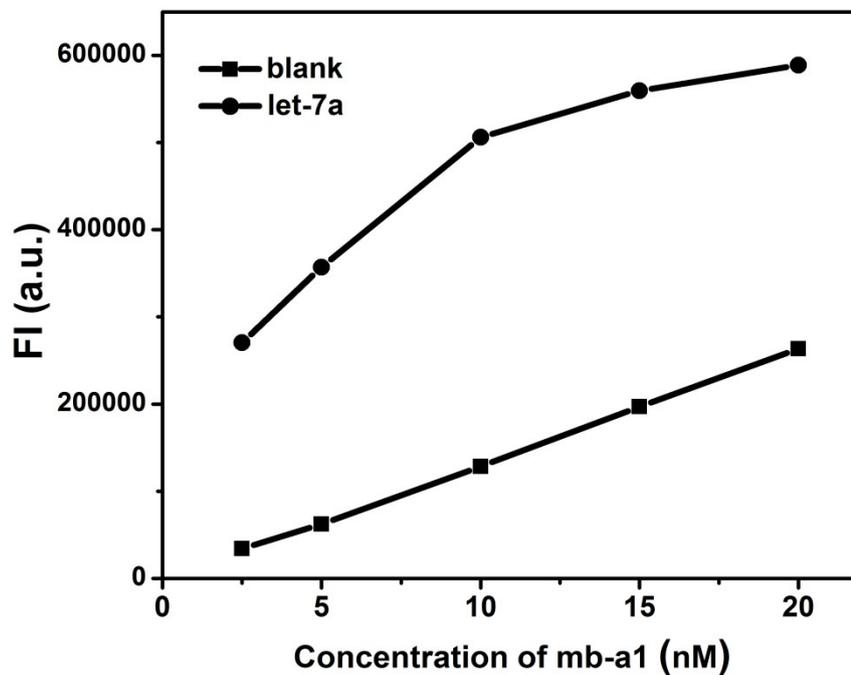


Fig. S4 The effect of the concentration of MB probe. The concentration of let-7a in the blank and the sample was 0 pM, 250 pM respectively. The reaction volume was 10 μ L and reaction products were diluted by 20 times with TE buffer solution before fluorescence measurement.

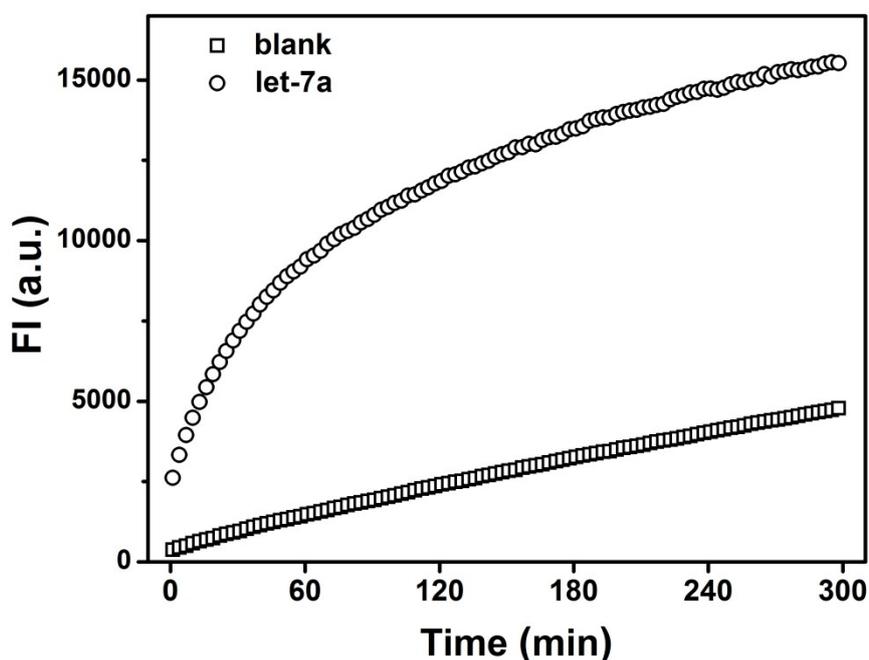


Fig. S5 The effect of the reaction time. The concentration of let-7a was 0 nM, 5 nM in the blank and the sample respectively.

Table S1 Comparison of different methods for miRNA detection

Method	Sensitivity	Selectivity	Simplicity	Time	ref
Northern Blotting with LNA probe	0.5 μ g total RNA	two-base difference	complex separation procedure	>17 h	1
RCA	10 fM	one-base difference	simple probe	8 h	2
Stem-loop RT-PCR	1 aM	one-base difference	complex probe design	>2 h	3
signal-amplifying ribozymes	5 nM	— ^a	complex probe design	10 min	4
DSN with DNA peroxidase probe	20 pM	— ^a	two-step detection procedure	2 h 45 min	5
DSN with DNA- AuNP probe	5 pM	— ^a	preparation of AuNP and conjugation of DNA- AuNP	5 h	6
DSN with 2-OMe-RNA MB probe	0.5 pM	one-base difference	2-OMe-RNA modified probe	40 min	7
this work	5 pM	one-base difference	Simple probe design and procedure	2 h	

^a “—” represents the data are not available.

References

1. E. Várallyay, J. Burgyán and Z. Havelda, *Nat. Protoc.*, 2008, **3**, 190-196.
2. Y. Cheng, X. Zhang, Z. Li, X. Jiao, Y. Wang and Y. Zhang, *Angew. Chem.*, 2009, **121**, 3318-3322.
3. C. Chen, D. A. Ridzon, A. J. Broomer, Z. Zhou, D. H. Lee, J. T. Nguyen, M. Barbisin, N. L. Xu, V. R. Mahuvakar and M. R. Andersen, *Nucleic Acids Res.*, 2005, **33**, e179.
4. J. S. Hartig, I. Grüne, S. H. Najafi-Shoushtari and M. Famulok, *J. Am. Chem. Soc.*, 2004, **126**, 722-723.

5. T. Tian, H. Xiao, Z. Zhang, Y. Long, S. Peng, S. Wang, X. Zhou, S. Liu and X. Zhou, *Chem. Eur. J.*, 2013, **19**, 92-95.
6. F. Degliangeli, P. Kshirsagar, V. Brunetti, P. P. Pompa and R. Fiammengo, *J. Am. Chem. Soc.*, 2014, **136**, 2264-2267.
7. X. Lin, C. Zhang, Y. Huang, Z. Zhu, X. Chen and C. James Yang, *Chem. Commun.*, 2013, **49**, 7243-7245.