

Electronic supplementary information

MnO₂ nanosheets-mediated CRISPR/Cas12a for detection of organophosphorus pesticide in environment water†

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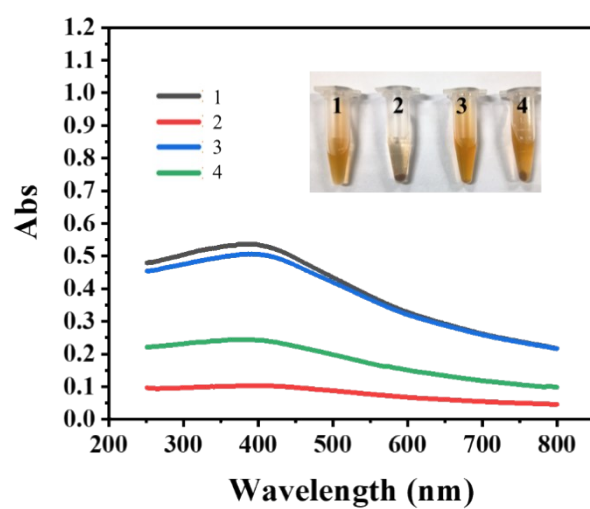


Fig. S1. The UV-vis spectra of MnO₂ nanosheets under different conditions: (1), MnO₂ nanosheets in ultra-pure water; (2), MnO₂ nanosheets in NaCl solution; (3), MnO₂-activator in NaCl solution; (4), MnO₂ nanosheets + activator in NaCl solution. Insert is the corresponding photograph.

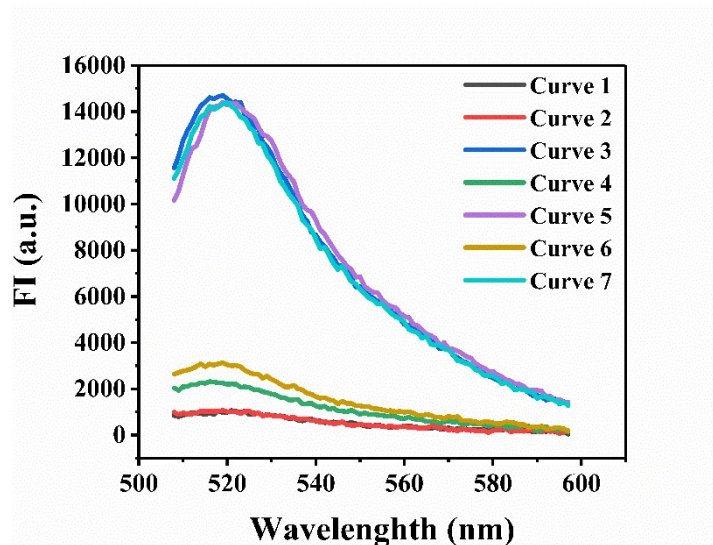


Fig. S2. The fluorescence spectra of different samples: Curve 1, FQ reporter; Curve 2, Cas12a/crRNA + FQ reporter; Curve 3, Activator + Cas12a/crRNA + FQ reporter; Curve 4, MnO₂-activator + Cas12a/crRNA + FQ reporter; Curve 5, AChE + ATCh + MnO₂-activator + Cas12a/crRNA + FQ reporter; Curve 6, AChE + ATCh + MnO₂-activator + Cas12a/crRNA + FQ reporter + DDVP; Curve 7, Activator + Cas12a/crRNA + FQ reporter + DDVP. DDVP: 50 µg mL⁻¹.

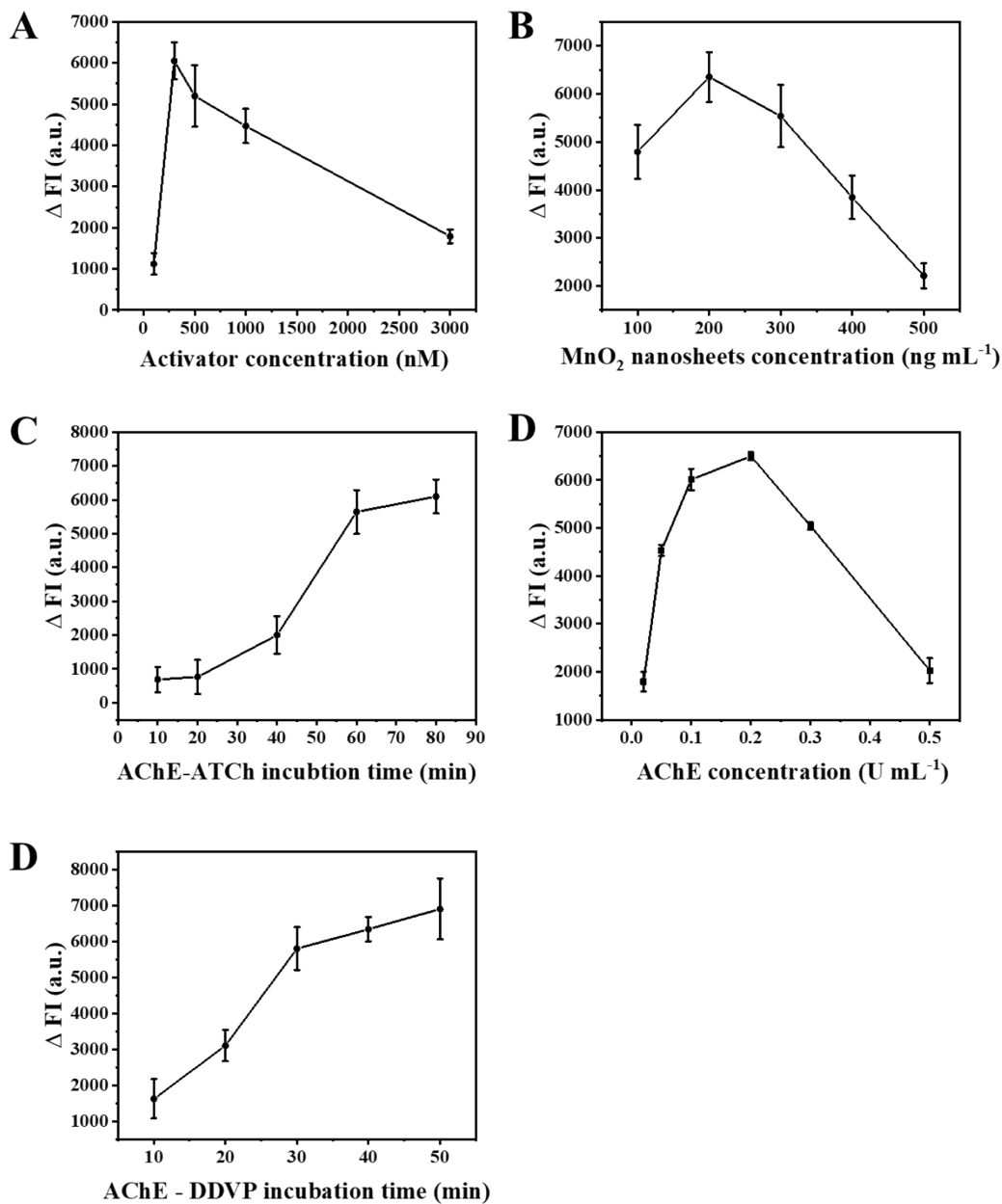


Fig. S3. (A) Effect of activator concentration on the sensing performance; (B) Effect of MnO_2 nanosheets concentration on the sensing performance; (C) Effect of AChE-ATCh incubation time on the sensing performance; (D) Effect of AChE concentration on the sensing performance; (E) Effect of AChE-DDVP incubation time on the sensing performance.

Table S1. Sequences of oligonucleotides used in this study.

Name	Sequence (5'-3')
Activator	GCT TAG AGT ATA GTA GTT GAT CG
crRNA	UAA UUU CUA CUA AGU GUA GAU AUC AAC UAC UAU ACU CUA A
FQ reporter	FAM-TTATT-BHQ1
Non-specific ssDNA for FAGE	CGC ACC TCG GAA TGT CGC GCA TGG TGC GCA

Table S2. Comparison between this method and other MnO₂ nanosheets-mediated sensors for DDVP detection.

Detect method	Linear range	LOD	References
Colorimetry	7–600 ng mL ⁻¹	2.3 ng mL ⁻¹	[1]
Colorimetry	5–60 μM	0.65 μM	[2]
Colorimetry	0.25–1.0 mM	42.94 μM	[3]
Fluorescence	10–1000 ng mL ⁻¹	3 ng mL ⁻¹	[4]
Fluorescence	137–2740 ng mL ⁻¹	0.406 ng mL ⁻¹	[5]
Fluorescence	5–1000 ng mL ⁻¹	0.135 ng mL ⁻¹	This work

References

1. J. Cao, M. Wang, Y. She, A.M. Abd El-Aty, A. Hacımüftüoğlu, J. Wang, M. Yan, S. Hong, S. Lao, Y. Wang, *Microchim. Acta*, 2019, **186**, 390.
2. W.S.T. Tun, C. Talodthaisong, S. Daduang, J. Daduang, K. Rongchai, R. Patramanon, S. Kulchat, *Mater. Chem. Front.*, 2022, **6**, 1487–1498.
3. S. L. D'souza, R. K. Pati, S. K. Kailasa, *Anal. Methods*, 2014, **6**, 9007–9014.
4. B. Lin, Y. Yan, M. Guo, Y. Cao, Y. Yu, T. Zhang, Y. Huang, D. Wu, *Food Chem.*, 2018, **245**, 1176–1182.
5. R. Fu, Y. Wang, Y. Liu, H. Liu, Q. Zhao, Y. Zhang, C. Wang, Z. Li, B. Jiao, Y. He, *Food Chem.*, 2022, **387**, 132919.