

Supporting Information

MoS₂ phase transition-assisted sensitive detection of cholesterol on the Copper oxide@Molybdenum disulfide nanocomposite

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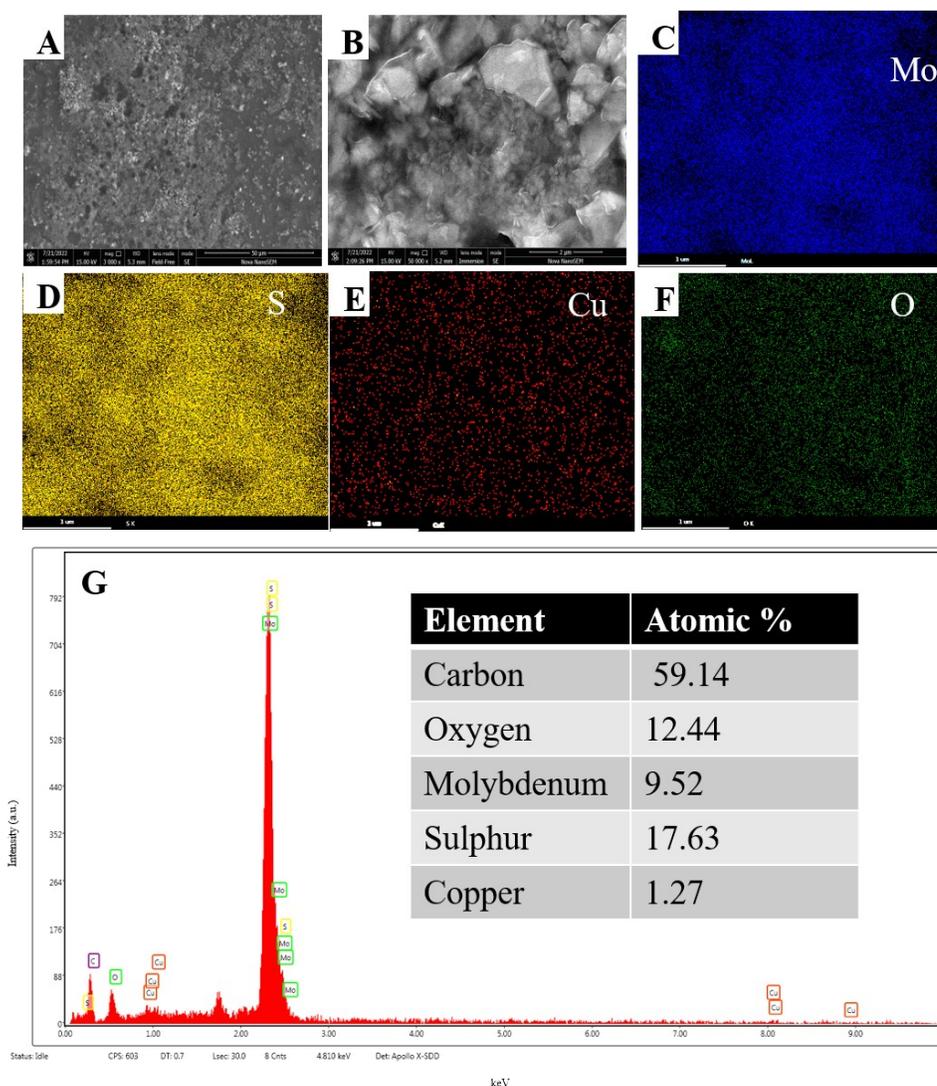


Fig. S1. (A) SEM image of CuO@MoS₂, (B) SEM field image for mapping, (C, D, E, and F) EDS mapping images for the elements Mo, S, Cu, and O. (G) EDS spectra of CuO@MoS₂.

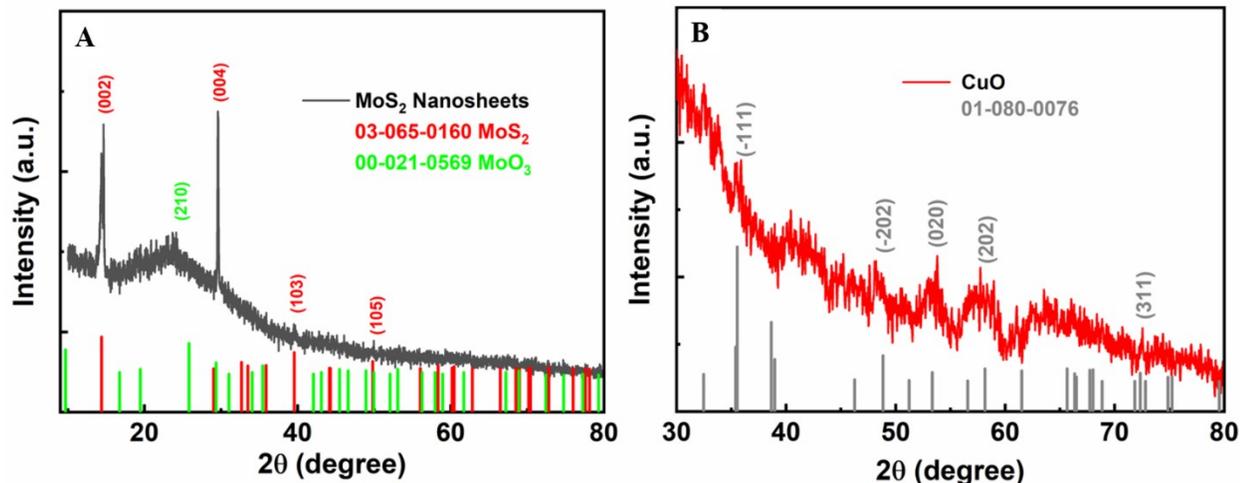


Fig. S2. XRD patterns and corresponding JCPDS data for (A) MoS₂ nanosheets and (B) CuO nanoparticles.

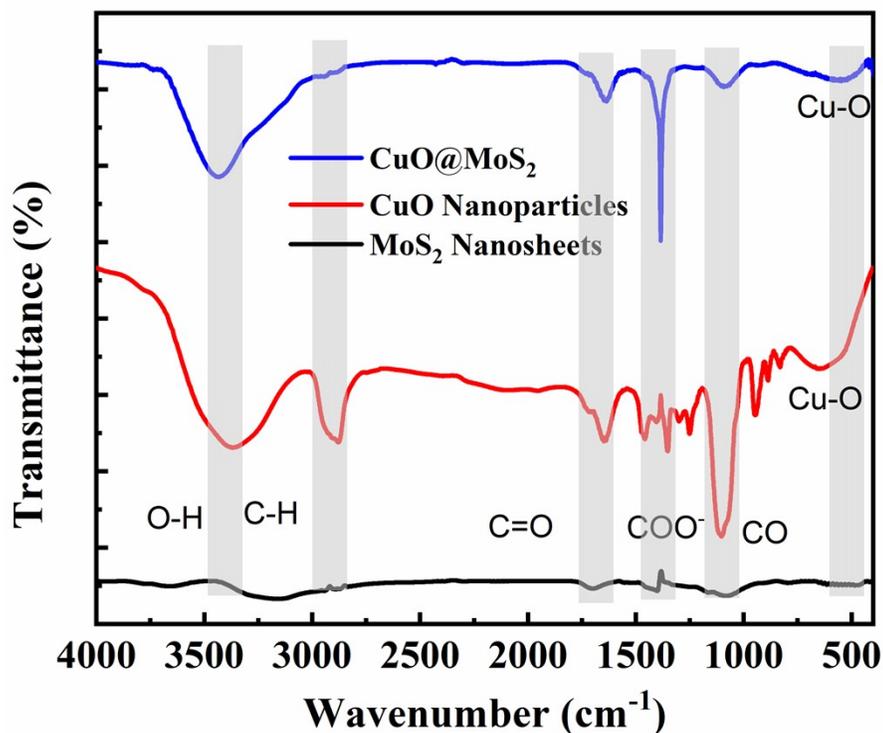


Fig. S3. FT-IR spectra of MoS₂ nanosheets, CuO nanoparticles, and CuO@MoS₂.

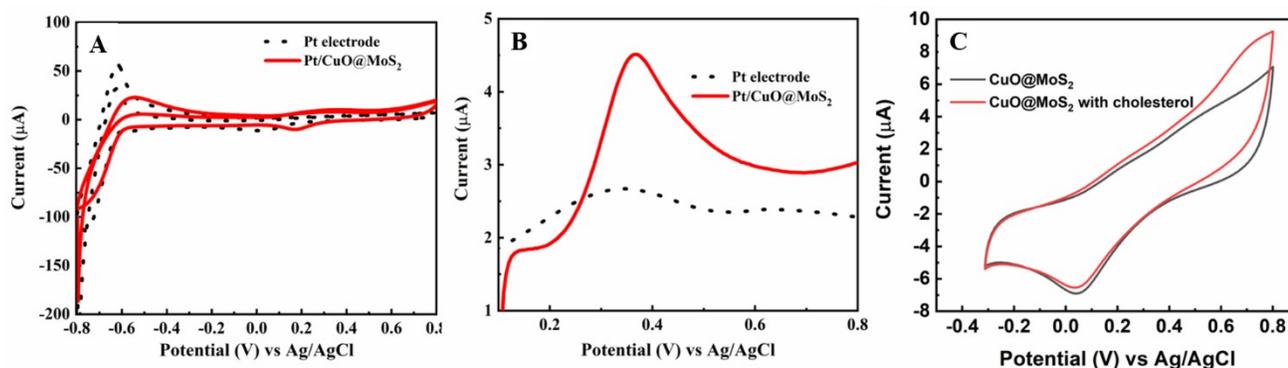
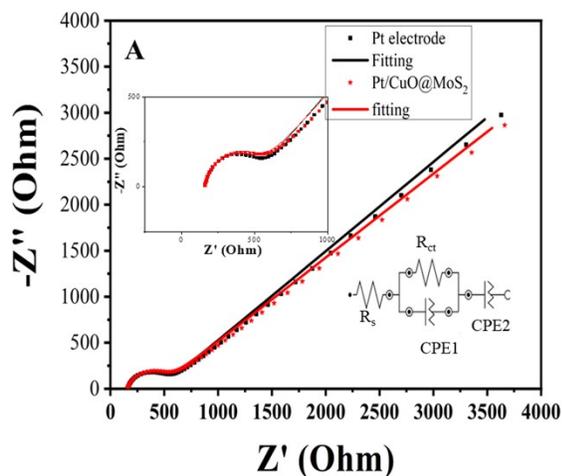


Fig. S4: (A) CV of Pt electrode and CuO@MoS₂, (B) DPV of Pt electrode and CuO@MoS₂, and (C) CV of CuO@MoS₂ with and without cholesterol in 0.1 M PBS.



B

Electrode	Rs (Ω)	Rct (Ω)	N of CPE 1	N of CPE 2
Pt	154	305	0.939	0.49
Pt/CuO@MoS ₂	151	287	0.95	0.471

Fig. S5: (A) Nyquist plot of (EIS) of Pt electrode and Pt/CuO@MoS₂ in 5×10^{-3} mol L⁻¹ K₃[Fe(CN)₆] and 0.1 mol L⁻¹ KCl. (B) Equivalent circuit element values corresponding to the fitting of the EIS data of the Pt electrode and its modification with CuO@MoS₂.

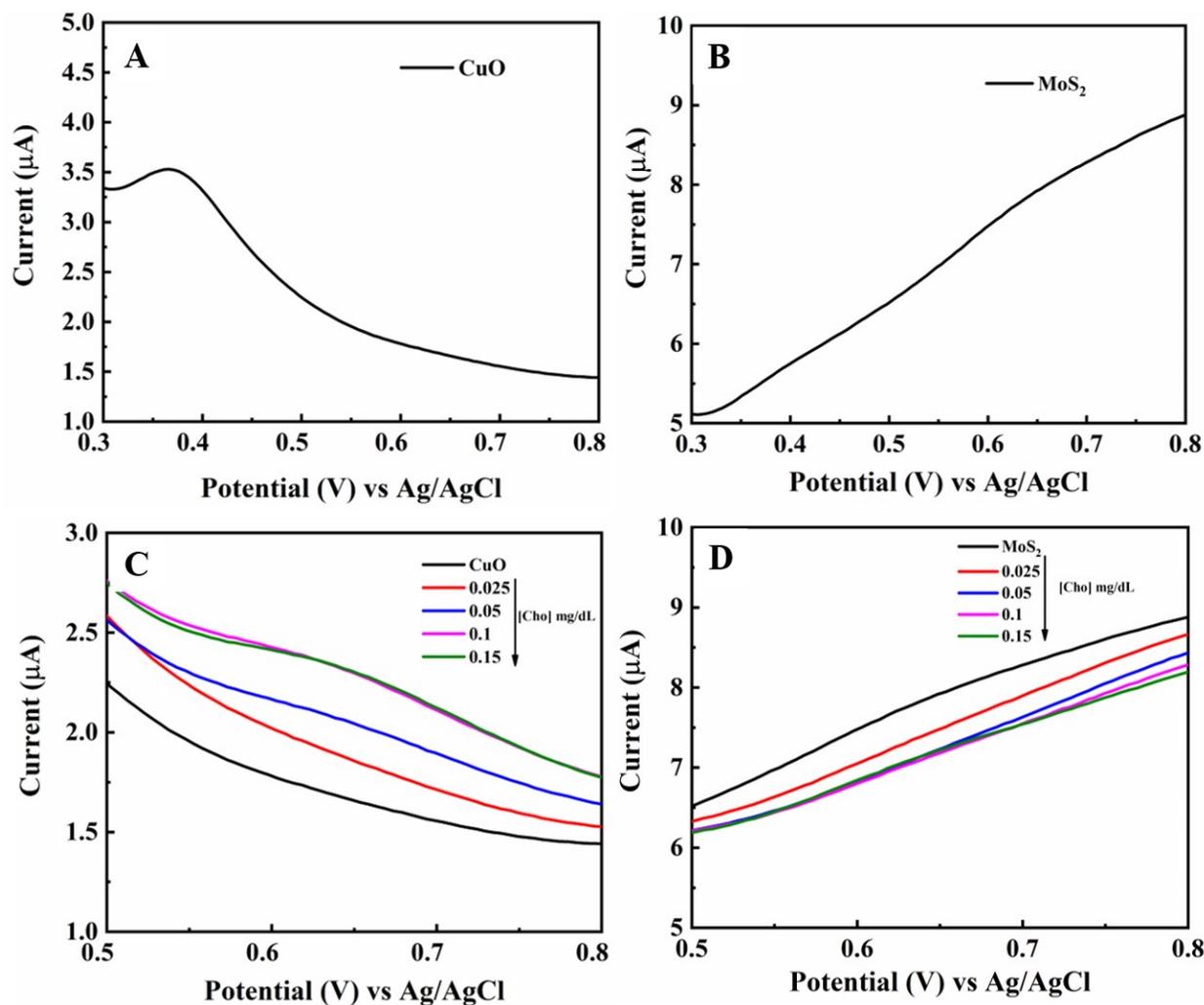


Fig. S6. Electrochemical response of (A) CuO, and (B) MoS₂ modified electrode in the potential window of 0.3 to 0.8 V in PBS. Effect of cholesterol addition on the Pt electrode modified with (C) CuO and (D) MoS₂.

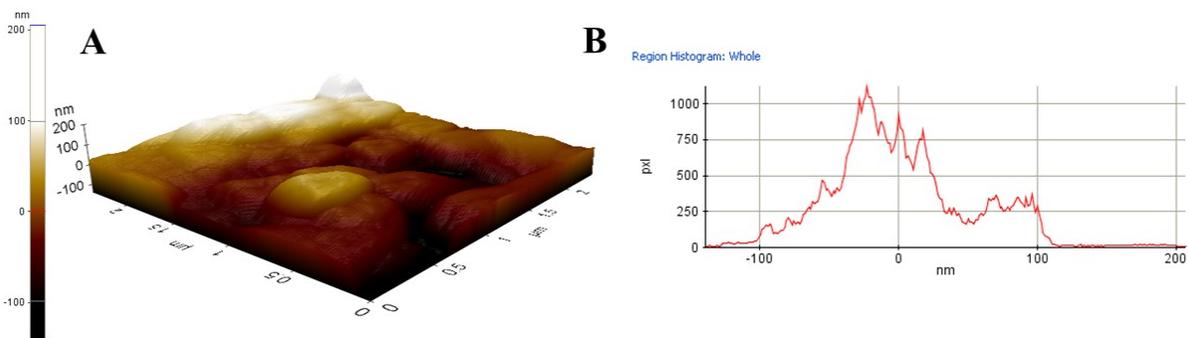


Fig. S7. AFM image of CuO@MoS₂ after cholesterol detection (A) Topography (B) Corresponding height profiles.

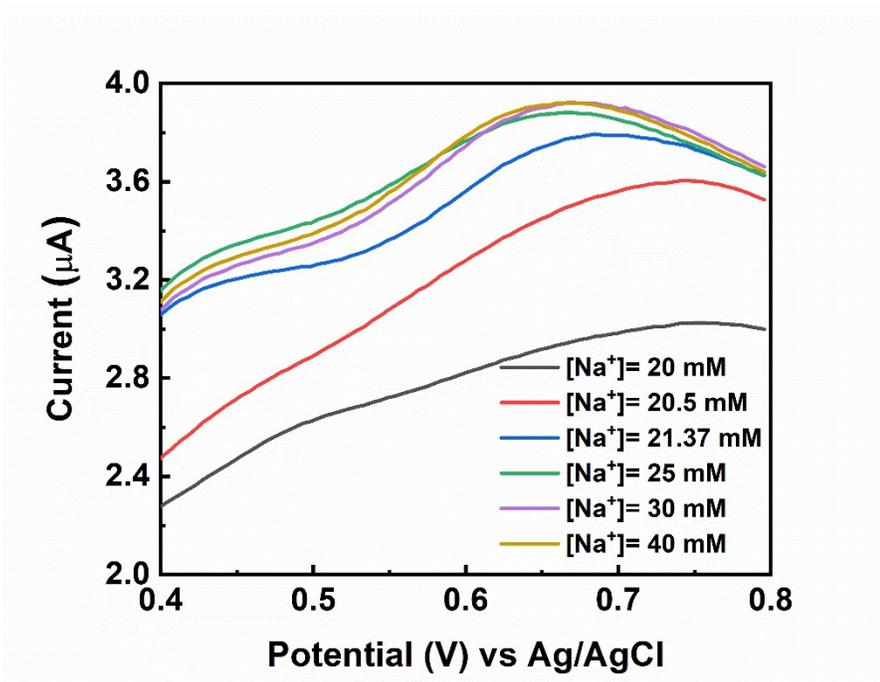


Fig. S8. The DPV response of 0.3 mg/dL cholesterol on the CuO@MoS₂ sensor in varying [Na⁺] buffer solutions.

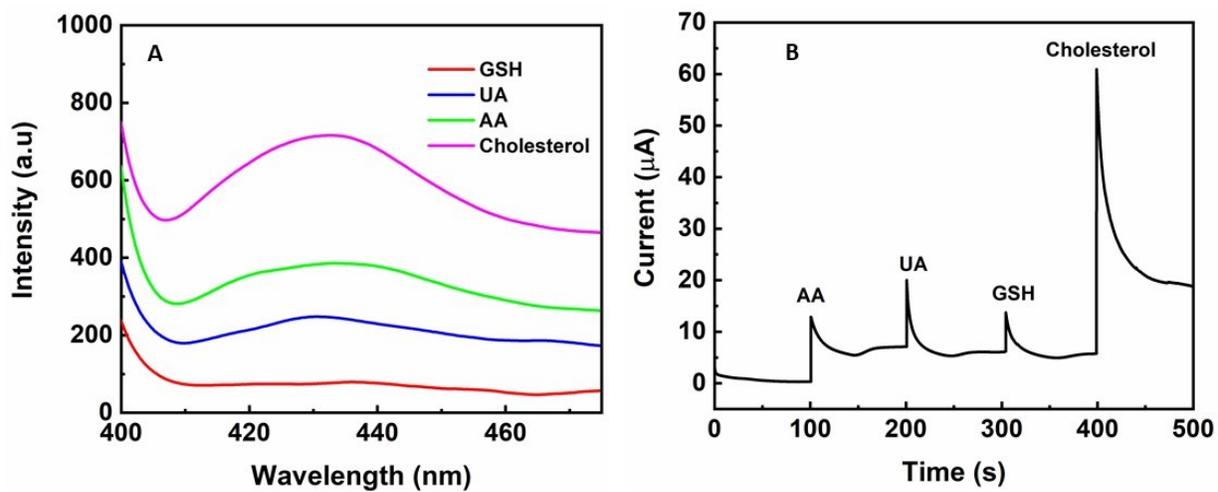


Fig.S9.(A). Fluorescence spectra of the electrolyte after Glutathione, AA, UA, and cholesterol detection (B) CA of AA, UA, Glutathione, and cholesterol with CuO@MoS₂ sensor .

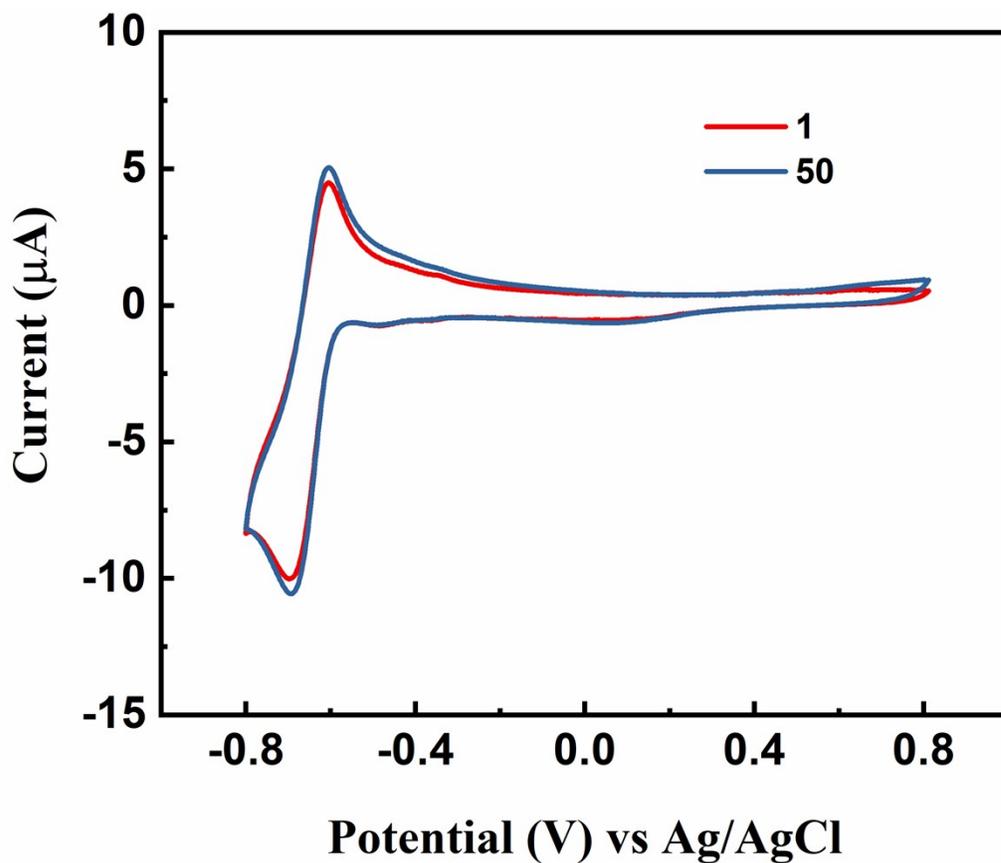


Fig. S10: CV of the first and fiftieth scans of Pt/CuO@MoS₂ in 0.1 M PBS.

Table. S1. Comparison of the CuO@MoS₂ sensor with reported cholesterol detection methods.

Sl No.	Detection Method	Material Used	Limit of Detection (LOD)	Detection Range	Reference
1	Electrochemical (Amperometric i-t method)	MoS ₂ -AuNPs	0.26 ± 0.015 μM	0.5–48 μM	1
2	Chemiluminescence	MoS ₂ and graphene quantum dots	35 nM	0.08–300 μM	2
3	Electrochemical (Amperometric i-t method)	MoS ₂ -Cu ₂ O hybrid nanoparticles	0.074 μM	0.25–25 μM	3
4	Electrochemical (Differential Pulse Voltammetry)	Poly(methyl orange)-modified NiO/MoS ₂ on screen-printed carbon electrode	0.24 mg/dL (6.2 μM)	1–15 mg/dL (25.86–387.9 μM)	4
5	Colorimetric	Au nanoparticles supported on MoS ₂ nanoribbons	0.015 mM	0.04–1 mM	5
6	Fluorometric	Ch Ox in MOF and Ag nanocluster @ MoS ₂ nanosheets	0.03 μM	0.06–15 μM	6
7	Electrochemical (Chronoamperometry)	ChOx/Pt wire based 3-D printed microneedle	0.5 μM	1 to 20 μM	7
8	Electrochemical (Differential Pulse Voltammetry)	CuO@ MoS ₂	679 nM (26.25x10 ⁻⁴ mg/dL)	1.29 μM-10.34 μM (0.05-0.4 mg/dL)	This work

References

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