

Electronic supplementary information

Smartphone-assisted colorimetric determination of uranyl ions in aqueous solutions

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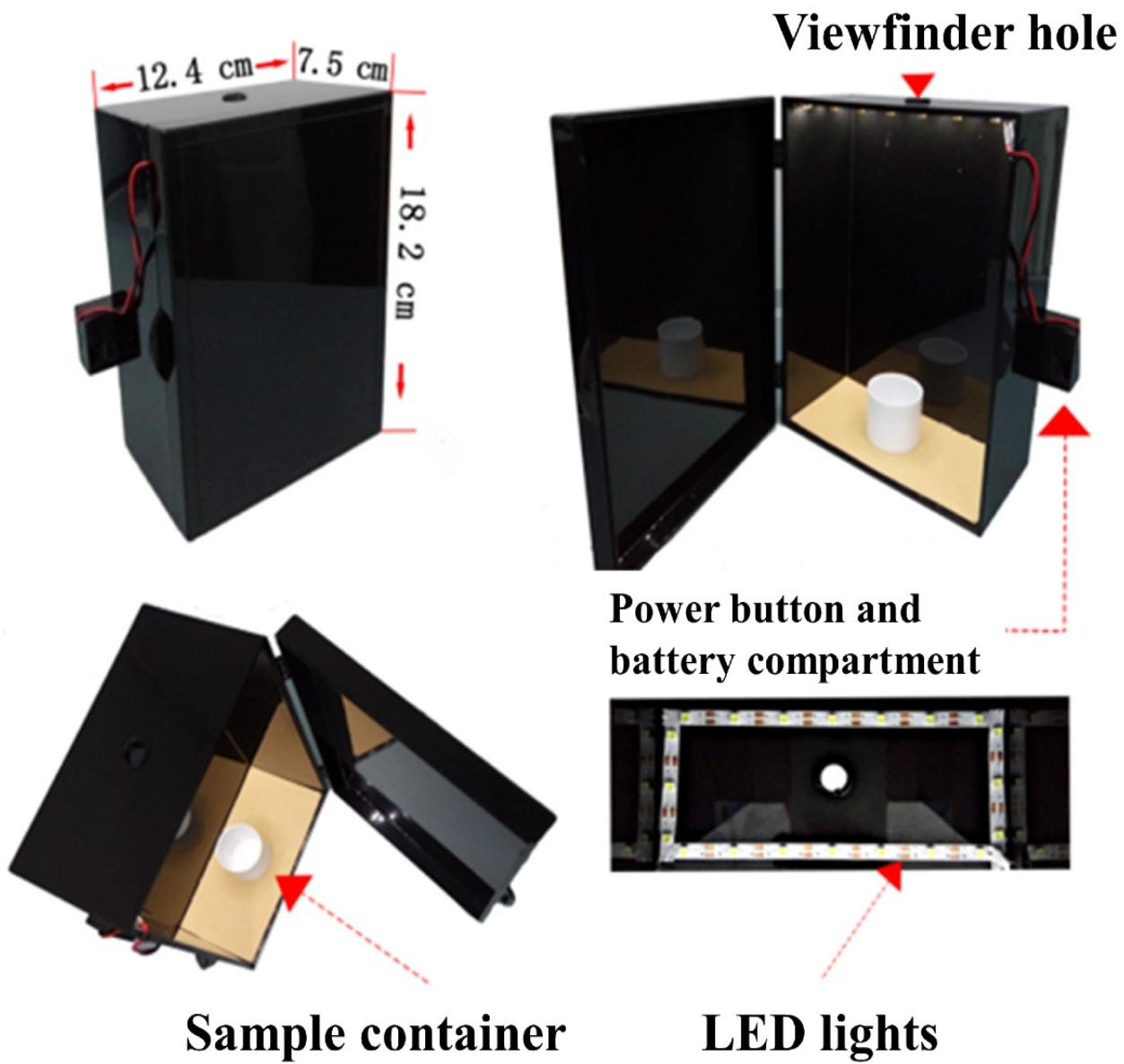


Fig. S1 Structure of the colorimetric viewfinder.

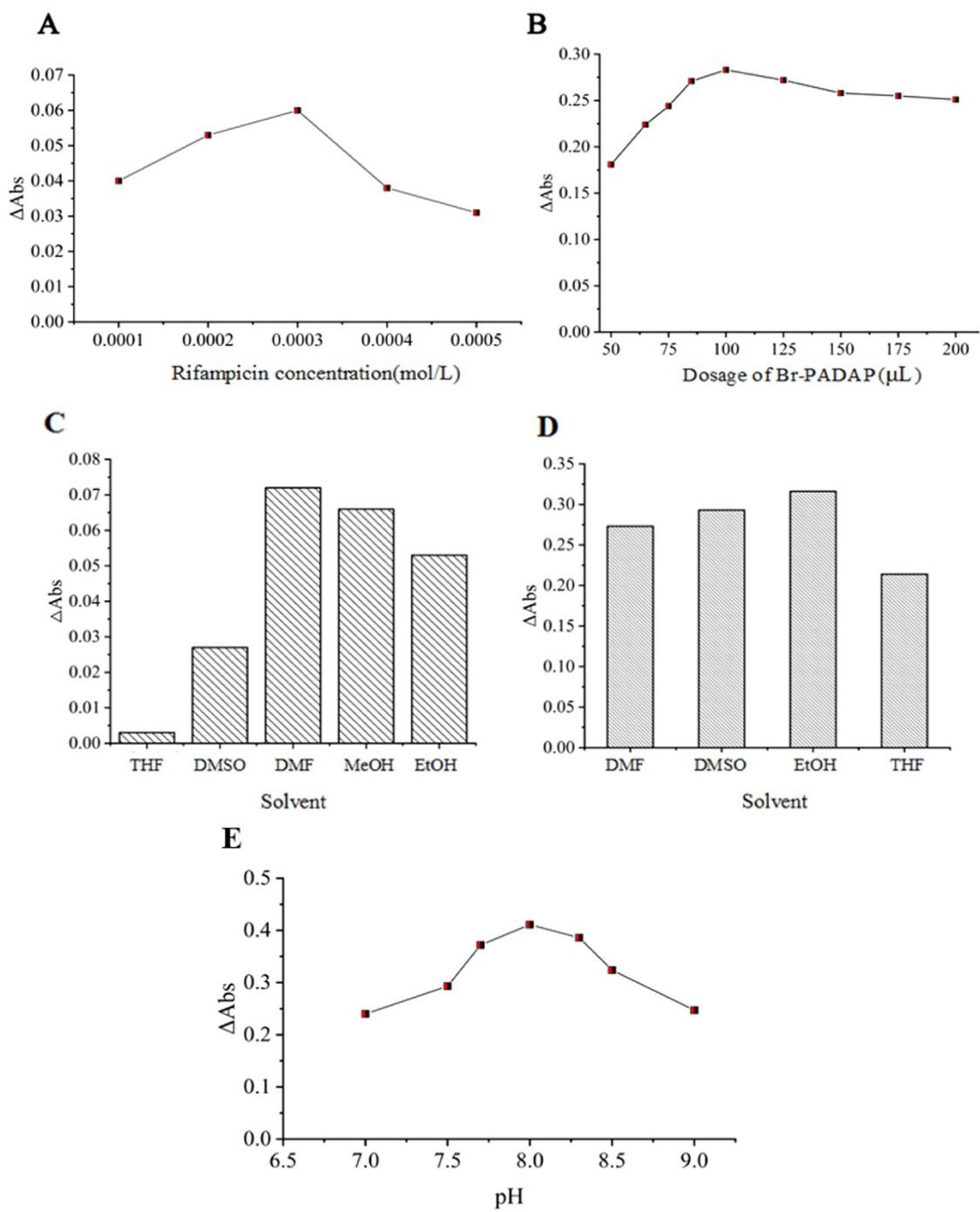


Fig. S2 Absorbance difference values of the rifampicin system with different concentrations of Rifampicin fixed at 2 mL dosage (A). Absorbance difference values of Br-PADAP at a fixed concentration of 1mM with different dosages (B). Absorbance difference values of rifampicin system(C) and Br-PADAP system (D) in different solvents. Absorbance difference values of Br-PADAP system in different pH values(E).

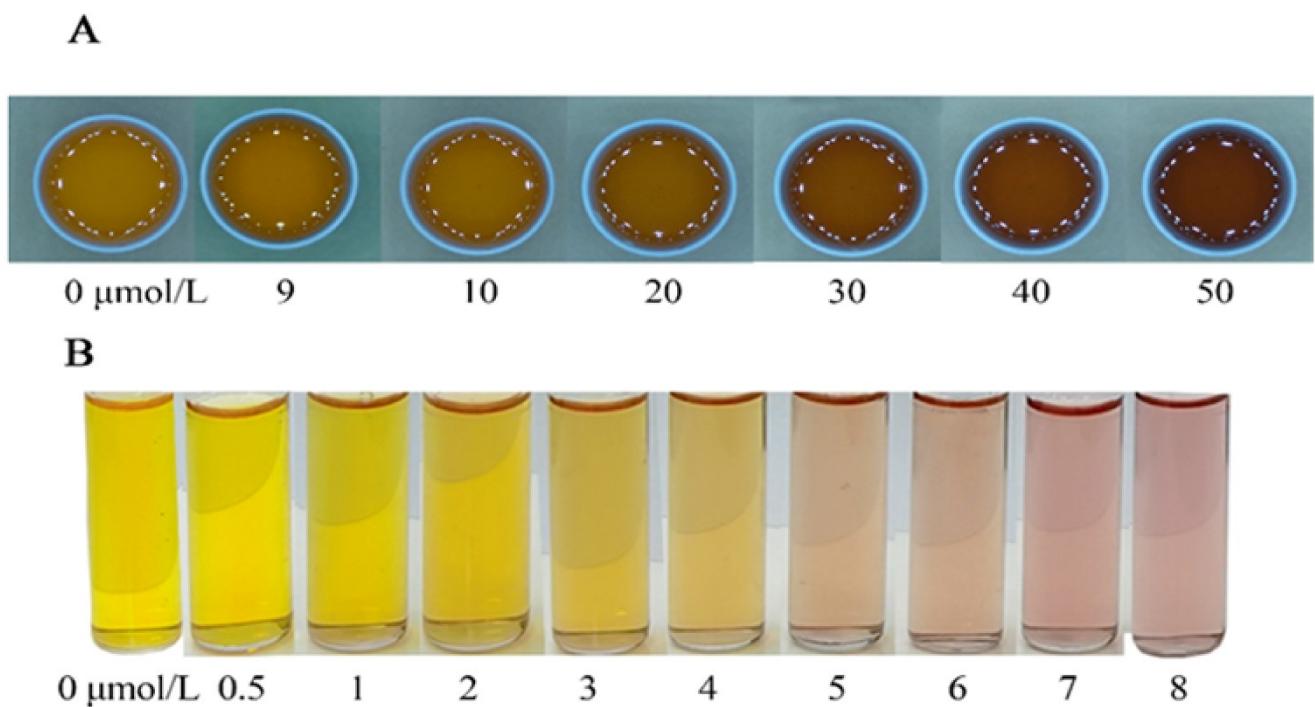


Fig. S3 Photographs of the detection solution of rifampicin system (A) and Br-PADAP (B) system mixed with different concentrations of UO_2^{2+} .

Table S1 Comparison of methods for determination of UO_2^{2+} .

Detection platforms	Method	Linear range	Detection limit	Ref.
Triton X-100 micelles-capped curcumin	colorimetry	3.7–14 μM	3.7 μM	¹
Nitrophenyldiacetic acids–AuNPs	colorimetry	0.5–3 μM	2 μM	²
VPA-AuNPs	colorimetry	0.5–10 μM , 4–20 μM	1.07 μM	³
Dual-colour label-free carbon dots	fluorescence	0–30.0 μM	8.15 μM	⁴
Europium metal-organic framework	fluorescence	12.5–87.5 μM	2.5 μM	⁵
Fe_3S_4 nanoparticles wrapped in a g-C ₃ N ₄ matrix	electrochemistry	0.05–8 μM	0.22 μM	⁶
Rifampicin system; Br-PADAP system	colorimetry	4–50 μM ; 0.9–7 μM	3.17 μM ; 0.89 μM	This work

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