

Supporting

Water-stable perovskite-on-polymer fluorescent microspheres for the simultaneous monitoring of pH, urea and urease

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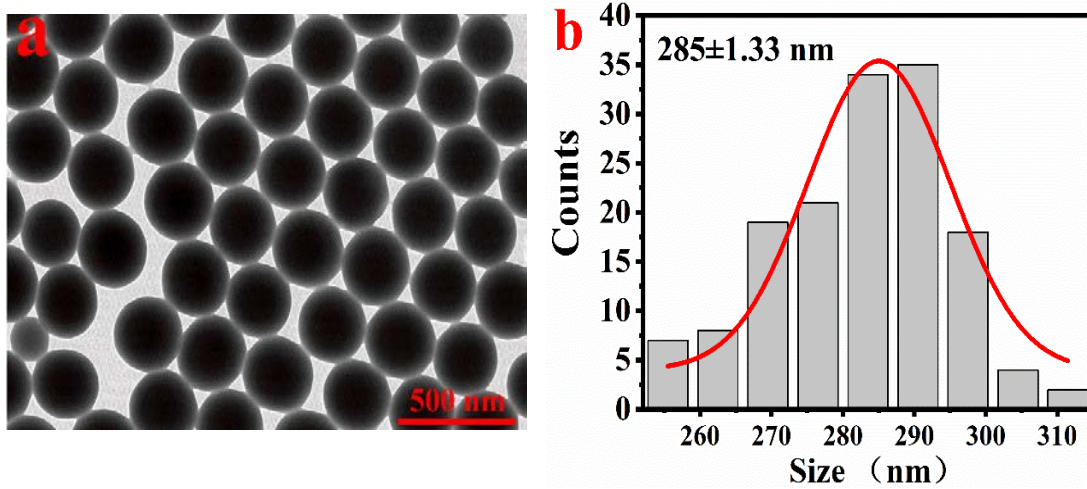


Fig. S1 (a) TEM image of polymer of polystyrene and poly (acrylic acid) (PS-PAA),
 (b) Corresponding size distribution of PS-PAA.

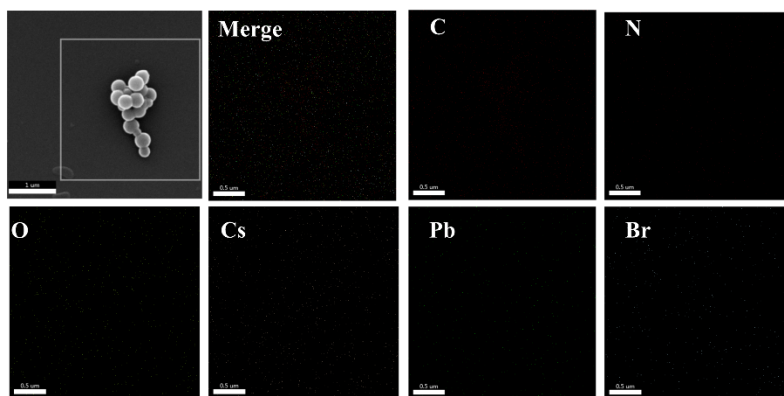


Fig. S2 SEM/EDX elemental mapping analysis of the synthesized PDPS composites.

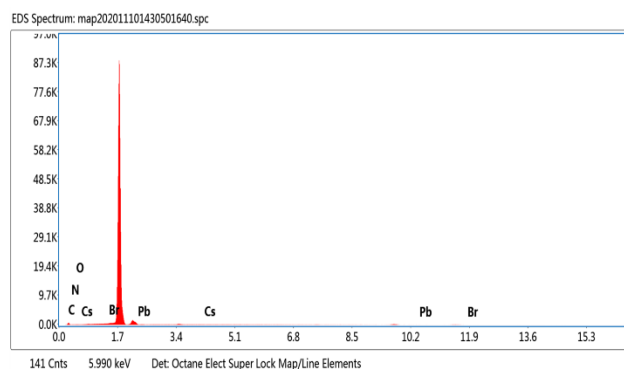


Fig. S3 The corresponding EDX images of PDPS composites.

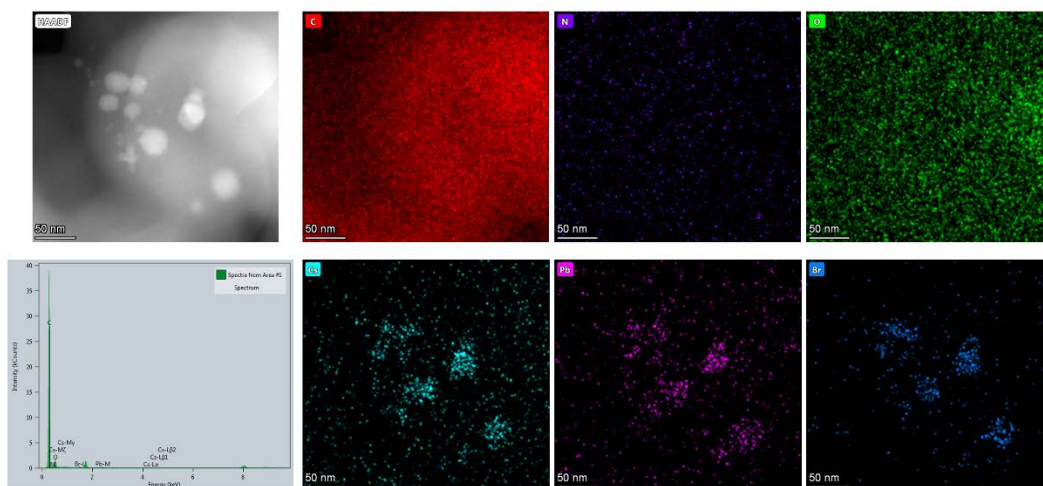


Fig. S4 The HAADF-STEM image and corresponding EDS layered images of the PDPS composites.

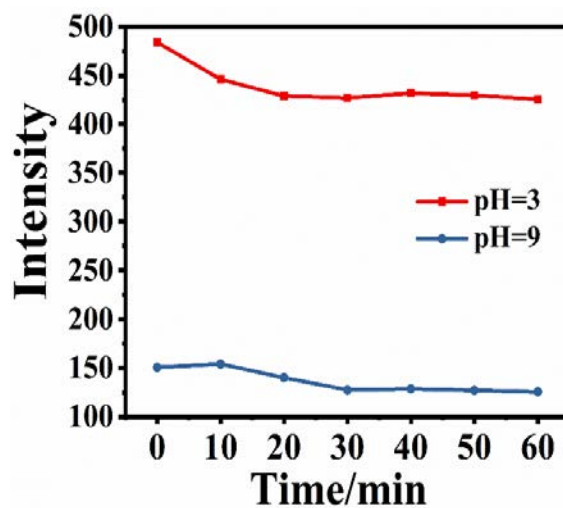


Fig. S5 Fluorescent intensity ($\lambda_{ex}=300$ nm) of the PDPS composites with different reaction time.

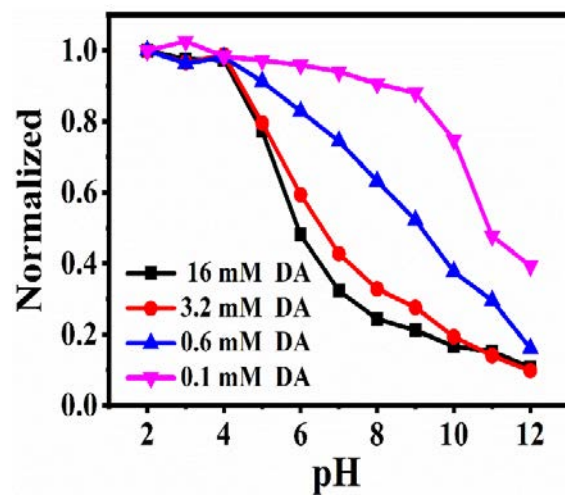


Fig. S6 The relationship between fluorescent intensity of the PDPS composites and pH at different concentration of DA

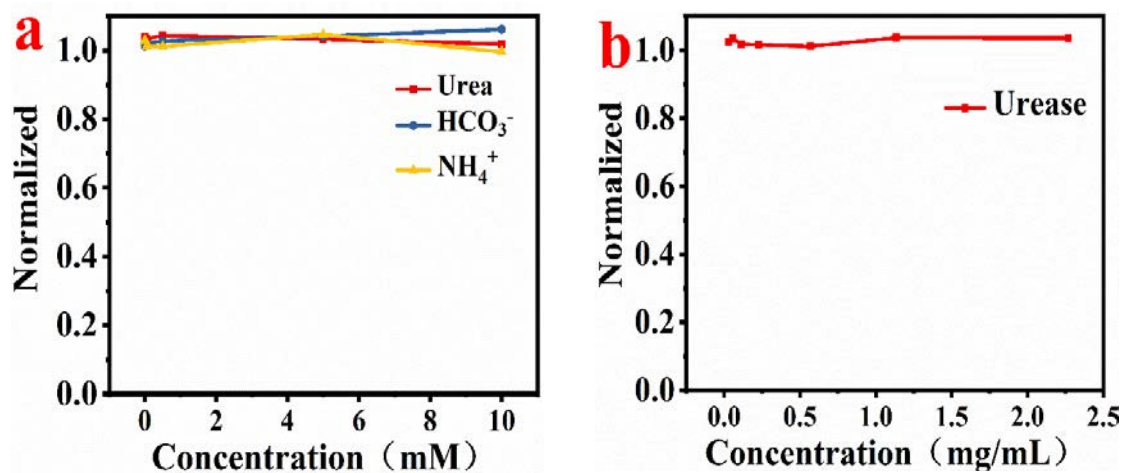


Fig. S7 The fluorescent intensity of the PDPS composites at different concentration of (a) Urea, HCO₃⁻, NH₄⁺, (b) Urease.

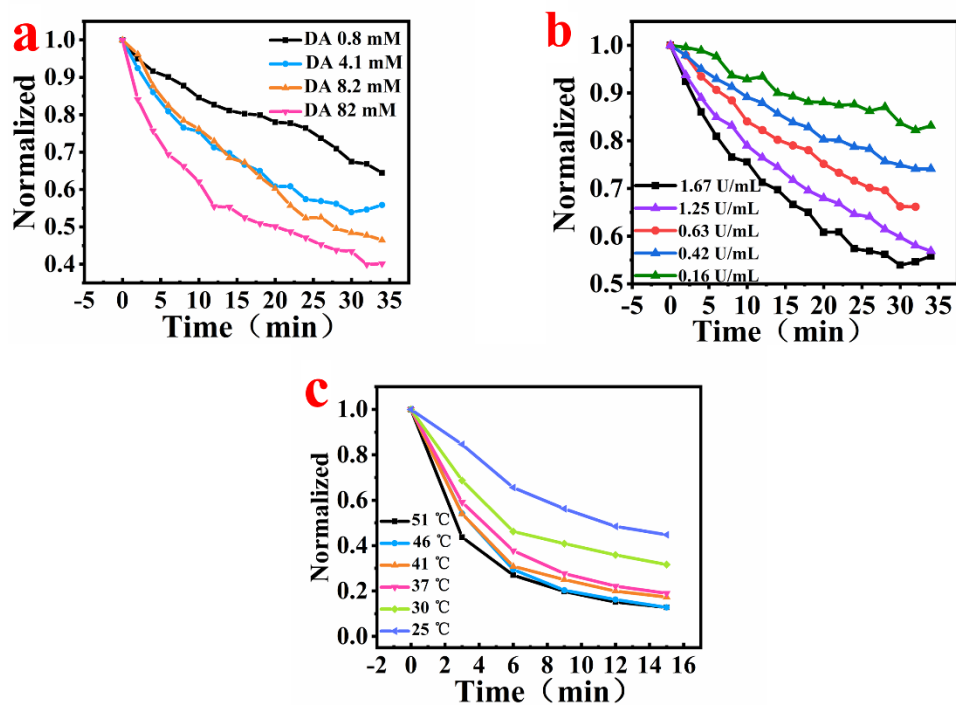


Fig. S8 The relationship between fluorescent intensity of the PDPS composites and the reaction time at different concentration of (a) DA, (b) Urease, (c) Temperature.

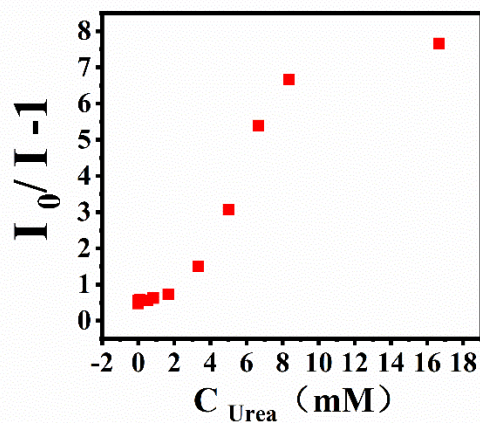


Fig. S9 The linear relationship between $(I_0/I-1)$ and urea concentration.

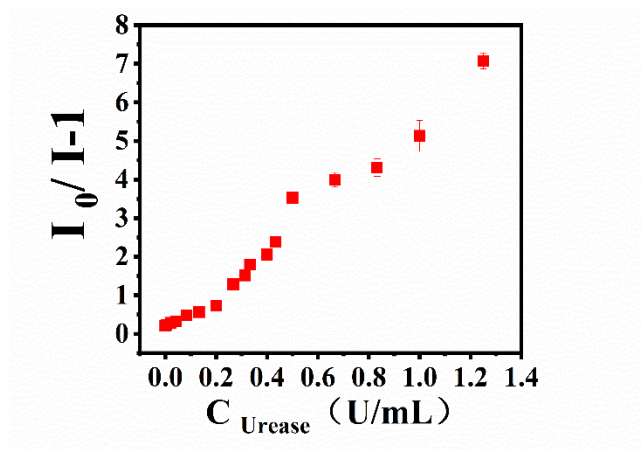


Fig. S10 The linear relationship between ($I_0/I-1$) and urease concentration.

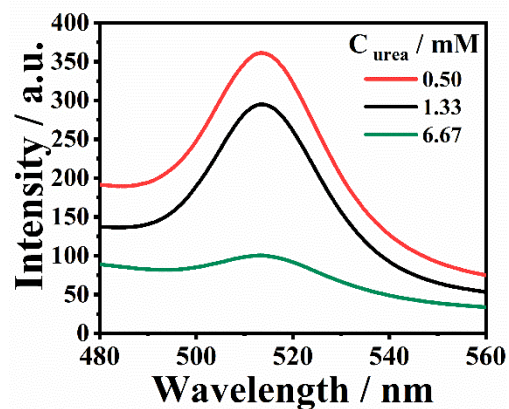


Fig.S11 Fluorescent emission spectra ($\lambda_{\text{ex}}=300 \text{ nm}$) of the PDPS composites with different concentrations of urea in real human serum samples.

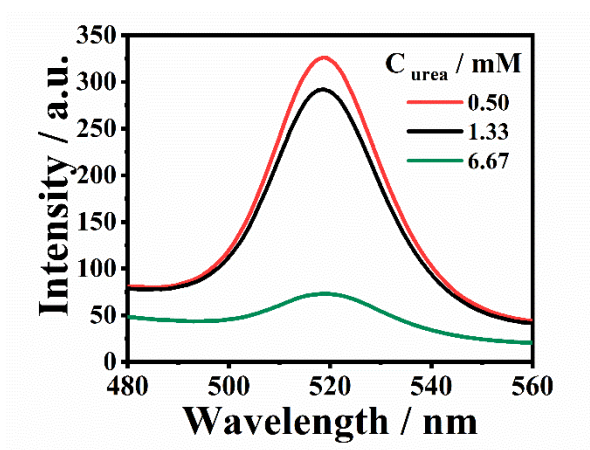


Fig.S12 Fluorescent emission spectra ($\lambda_{\text{ex}}=300 \text{ nm}$) of the PDPS composites with different concentrations of urea in real human urine samples.

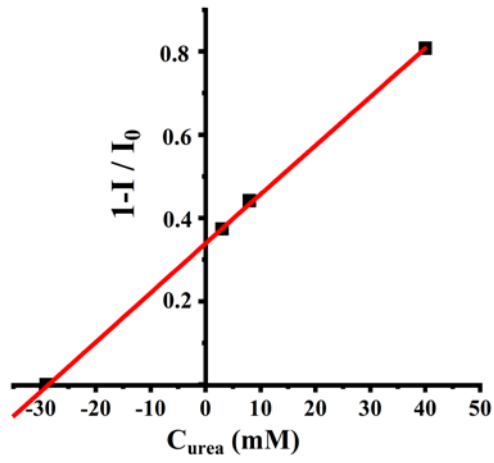


Fig.S13 Standard addition approach graph of detecting unknown concentrations of urea in real human serum samples.

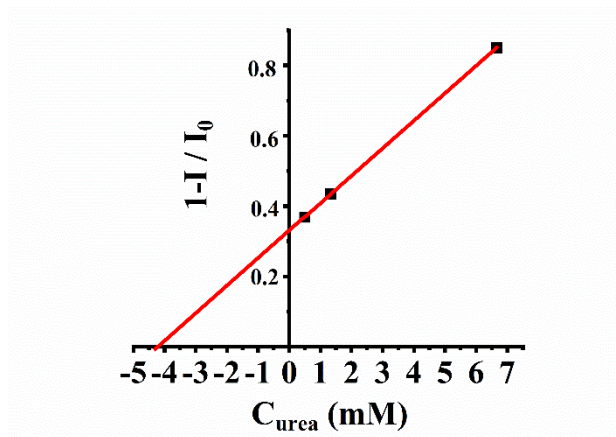


Fig.S14 Standard addition approach graph of detecting unknown concentrations of urea in real human serum samples.

Table S1

The comparison for urea detection among different sensors.

Type	Materials	Response time	Linear range	Detection limit	References

Electrochemistry	Yb ₂ O ₃	-	0.05-19 mM	2 μM	[4]
Fluorescence	P dots	45 min	0 -1 mM	0.02 mM	[5]
Fluorescence	CuO NPs	45 min	0.0375-0.3 mM	27 μM	[6]
Electrochemistry	MWCNT-PoT	-	0.1-11 mM	0.03 mM	[7]
Phosphorescence	MPA-CdTe quantum dots	10 min	0.016-0.16 mM	0.003 mM	[8]
Electrochemistry	urease/SF/aminated GCE	-	0.3-2.7 mM	0.163 mM	[9]
Fluorescence	MoS ₂ QDs	44 min	5-700 μM	1.8 μM	[10]
Fluorescence	PDPS composites	30 min	1.67 μM-6.67 mM	1.67 μM	This work

Table S2

The comparison of constructed sensor with other perovskite-based sensors.

Materials	Analyte	Signal	Stability	Detection Condition	References
CsPbBr ₃	Picric acid	Fluorescence	Vulnerable to humidity /pH/temperature	Cyclohexane	[11]
CsPbBr ₃ /cellulose composites	Chloride and Iodide Ions	Absorbance	Vulnerable to humidity /pH/temperature	Water	[12]
CH ₃ NH ₃ PbBr ₃	Hg ²⁺	Fluorescence	Vulnerable to humidity /pH/temperature	Toluene	[13]
CsPbBr ₃	Cu ²⁺	Fluorescence	Vulnerable to humidity /pH/temperature	Cyclohexane	[14]
(C ₆ H ₅ NH ₃) ₂ Pb ₃ I ₈ ·2H ₂ O	Fe ³⁺	Fluorescence	Vulnerable to humidity /pH/temperature	DMF	[15]
CsPbBr ₃ /MIP	Pesticide	Fluorescence	Vulnerable to humidity /pH/temperature	Dichloromethane	[16]
PDPS composites	pH/urea/urease	Fluorescence	Hyperstatic	Water	This work

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