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Supporting information for

Impact of nucleic acid encapsulated MOF crystal phase on protein corona formation

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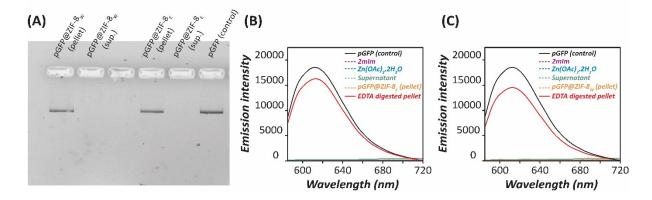


Fig S1. (A) Agarose gel electrophoresis image of the pGFP@ZIF-8_W, pGFP@ZIF-8_E pellet, supernatant and control pGFP. (B) Fluorescence spectroscopy of PI-added with the precursors, supernatant, pGFP@ZIF-8_E pellet and EDTA-digested pellet. (C) Fluorescence spectroscopy of PI-added with the precursors, supernatant, pGFP@ZIF-8_W pellet and EDTA-digested pellet.

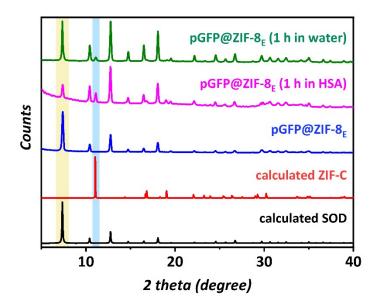


Fig S2. XRD of pGFP@ZIF-8_E in as-prepared condition, 1h HSA treatment, and 1 h water treatment.

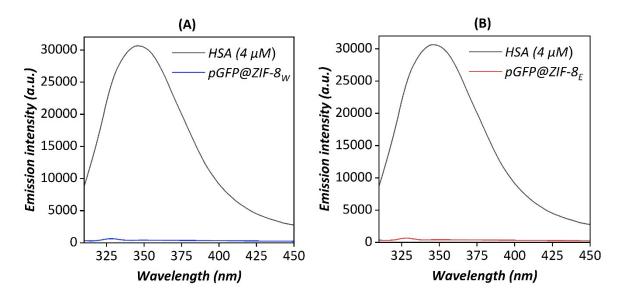


Fig S3. Fluorescence emission spectra of HSA molecules and untreated (A) pGFP@ZIF-8_W and (B) pGFP@ZIF-8_E.

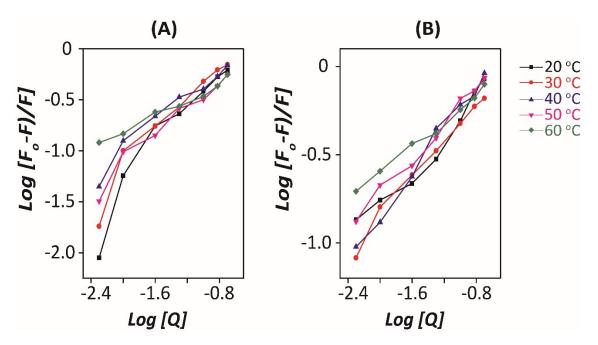


Fig S4. Double logarithmic regression plots of $log[(F_o-F)/F]$ vs. log[Q] for binding of HSA with (A) pGFP@ZIF-8_E and (B) pGFP@ZIF-8_W.

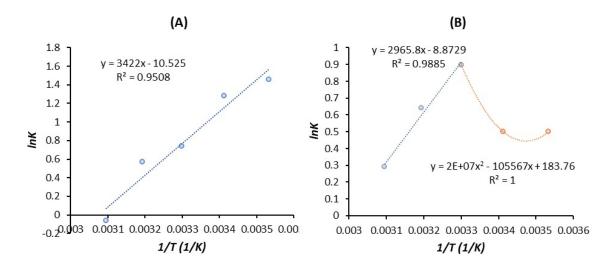


Fig S5. van't Hoff plot for (A) pGFP@ZIF- 8_E @HSA and (B) pGFP@ZIF- 8_W @HSA to determine interaction of ZIFs to HSA.

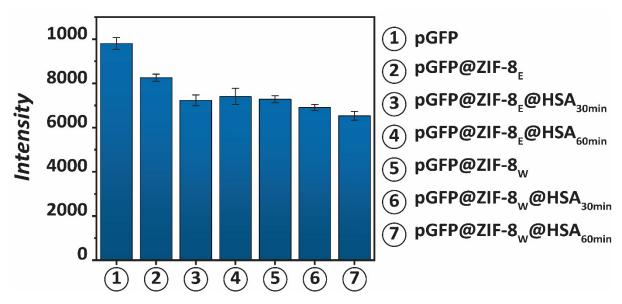


Fig S6. Protection of nucleic acid in the bio-composite after DNase treatment.

Table S1. Stern-Volmer constants and thermodynamic parameters for pGFP@ZIF- 8_E @HSA.

Temp (°C)	Ksv (M ⁻¹)	K (M ⁻¹)	n (M ⁻¹)	ΔH° (J/mol)	ΔS° (J/mol)	ΔG° (kJ/mol)
20	3.03	4.29	1.03	1.500	_	0.386
30	3.45	3.60	0.89	0.679		0.398
40	2.96	2.08	0.66	0.186	-1.311	0.411
50	2.47	1.76	0.70	-0.150	_	0.423
60	2.03	0.93	0.39	-0.946	_	0.436

 $\textbf{Table S2.} \ Stern-Volmer \ constants \ and \ thermodynamic \ parameters \ for \ pGFP@ZIF-8_W@HSA.$

Temp (°C)	Ksv (M ⁻¹)	K (M ⁻¹)	n (M ⁻¹)	ΔH° (J/mol)	ΔS° (J/mol)	ΔG° (kJ/mol)
20	3.73	1.65	0.50	-257.33 x 10 ³	-409.11	129.61 x 10 ³
30	2.84	1.65	0.54	-219.87 x 10 ³	-283.37	123.74 x 10 ³
40	3.99	2.45	0.62	-184.81 x 10 ³	-169.49	127.87 x 10 ³
50	3.66	1.90	0.50	1.176	-1.104	0.358
60	2.84	1.34	0.36	1.141	-	0.369