Secondary Structure-Induced Aggregation by Hydrogen Peroxide:

A Stimuli-Triggered Open/Close Implementation by Recombination

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Supporting Information Table of Contents

I. Experimental Details

- Chemicals and Materials
- Characterizations
- Synthesis of mono(4-(hydroxymethyl)phenylboronic acid pinacol ester) succinate (HMPBApin-SA)
- Synthesis of 4-hydroxy genistein 4,7- hydroxy -(4-(hydroxymethyl) phenylboronic acid pinacol ester) succinate ester (Ge-di(HMPBA-pin))
- Synthesis of TPGS-A
- Self-assembly of Ge-di(HMPBA-pin)-TPGS nanorods
- Au NRs were synthesized following the seeded growth method
- Citrate-stabilized Au NPs with an average diameter of ~13 nm synthesized according to literature procedures
- Synthesis of AIE monomer, TPE-CHO
- Synthesis of MoS₂ QDs
- Quantum Yields (QYs) Measurement (Table S1)

II. Supplemental Figures of Polymer Nanorods

Scheme S1. Synthesis route of a Ge-crosslinker (Ge -di(HMPBA-pin)).

Figure S1. ¹H NMR spectra of HMPBA-pin-SA and Ge -di(HMPBA-pin) in DMSO.

Figure S2. ESI-MS spectra of compound Ge -di(HMPBA-pin).

Figure S3. (a) TEM image of the nanorods, (b-d) HRTEM image of an individual nanorod created within 5 seconds. The inset shows the corresponding lattice fringes of the area within the dotted circle.

Figure S4. Schematic representation for the spherical micelles based on self-assembly of Gedi(HMPBA-pin) and TPGS-A.

Figure S5. ¹H NMR spectra of TPGS, TPGS-A.

Figure S6. ESI-MS spectra of TPGS (a) and TPGS-A (b).

Figure S7. Schematic representation for the synthesis of spherical micelles via self-assembly of Genistein and TPGS.

Figure S8. Schematic representation for the synthesis of nanoribbon via self-assembly of HMPBA-pin-SA and TPGS.

Figure S9. (a) UV-vis absorption spectra and (b) absorbance intensity ratios, A_{637}/A_{520} , recorded for the aqueous colorimetric assay (3 nM of AuNPs) and 1.5 µM polymer upon adding varying amount of H₂O₂ (0-150 equiv. relative to polymer moieties) at pH 7.4 and 25 °C. Optical visualization of the colorimetric assay (3 nM of AuNPs) and 1.5 µM polymer in the absence of H₂O₂ (left) and 1 h after the addition of the aqueous solution of H₂O₂ (100 equiv. relative to nanorod moieties) at pH 7.4 and 25 °C. (c, d) The TEM images obtained by drying the two aqueous dispersions corresponded to the insert in (a).

Figure S10. (a) HR TEM image of the aggregated MoS_2 QDs. (b) A partially destroyed micelle after the high-energy electron beam irradiation of 200 kV. The inset shows the corresponding lattice fringes of the area within the dotted circle. (c) Hydrodynamic diameter, D_h , distribution of dissociated the aggregated MoS_2 QDs in aqueous solution (0.5 mg/L).

Figure S11. (a) excitation-dependent PL behavior of MoS_2 QDs. (b) aggregated MoS_2 QDs under different excitation wavelengths.

Figure S12. ¹H NMR spectra of TPE-CHO in CDCl₃.

Figure S13. The digital photographs of TPE on the micelles surfaces in the faint yellow powder stored at room temperature (a); (b) the blue fluorescence of TPE on the micelles surfaces under UV light ($\lambda_{ex} = 365$ nm).

Figure S14. (a, b) The TEM images obtained by drying the two aqueous dispersions corresponded

to the inset in every picture at the same condition. (c) TEM image of the Cu²⁺ dispersion. (d) TEM image of the aggregated micelles.

Figure S15. Polymer nanorod glutamate toxicity-induced cell death. Concentration-dependent effect of water soluble nanorod on HT22 cell viability alone and with 4 mM glutamate. Nanorod was added to the cell media prior to the glutamate exposure and cell viability was estimated 24 h after addition. Nanorod alone upto 30 μ g had no effect on cell viability whereas glutamate exposure significantly reduced the survival of cells to nearly 30% of control. Pretreatment of cells with Nanorod resisted the effect of glutamate toxicity thereby showed significant reduction in cell mortality. Nanorod concentration of 10 μ g showed maximum protection to glutamate induced cell death in HT22 cells.

Figure S16. Polymer nanorod glutamate toxicity-induced cell death. Representative photomicrograph of HT22 cells treated with nanorod, glutamate, genistein and both. Data are the representation of 3 or more independent experiments conducted in triplicate. Glu=glutamate, Ge=genistein, Glu + naorod=glutamate + nanorod and Glu+nanorod-10 to 30 = glutamate (4 mM) + nanorod (10-30 µg).



Scheme S1. Synthesis route of Ge-crosslinker (Ge -di(HMPBA-pin)).



Figure S1. ¹H NMR spectra of HMPBA-pin-SA (a) and (b) Ge -di(HMPBA-pin) in DMSO.



Figure S2. ESI-MS spectrum of compound Ge -di(HMPBA-pin).



Figure S3. (a) TEM image of the nanorods, (b-d) HRTEM image of an individual nanorod created within 5 seconds. The inset shows the corresponding lattice fringes of the area within the dotted circle.



Figure S4. Schematic representation for the synthesis of spherical micelles via self-assembly of Ge-di(HMPBA-pin) and TPGS-A.



Figure S5. ¹H NMR spectra of TPGS (top), and TPGS-A (bottom).



Figure S6. ESI-MS spectra of (a) TPGS and (b)TPGS-A.



Figure S7. Schematic representation for the synthesis of spherical micelles via self-assembly of Genistein and TPGS.



Figure S8. Schematic representation for the synthesis of nanoribbon via self-assembly of HMPBA-pin-SA and TPGS.



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Figure S11. (a) excitation-dependent PL behavior of MoS_2 QDs. (b) aggregated MoS_2 QDs under different excitation wavelengths.

Quantum Yields (QYs) Measurement

Rhodamine B in ethanol (QY = 0.65, from literature¹) was selected as a control standard. The QYs of MoS_2 QDs and aggregated MoS_2 QDs were estimated according to the follow equation:

$$\Phi = \Phi_r \times \frac{I}{A} \times \frac{A_r}{I_r} \times \frac{n^2}{n_r^2}$$

Where the Φ is the QY, I is the integrated PL emission intensity (excited at 270 nm for MoS₂ QDs and at 285 nm for aggregated MoS₂ QDs, respectively), n is the refractive index (= 1.3614 for ethanol), and A is the absorbance value at the excitation wavelength of 270 nm and 285 nm in ethanol, respectively (the value of A should be less than 0.1 at the excitation wavelength). The subscript "r" refers to the standards.

Samples	Integrated emission intensity (I)	Abs. 270 nm (<i>A</i>)	Refractive index	QY
			of solvent (<i>n</i>)	(Φ)
Rhodamine B	20255	0.0195	1.362	0.65
MoS ₂ QDs	10125	0.0371	1.362	0.08
Samples	Integrated emission intensity (1)	Abs. 285 nm (<i>A</i>)	Refractive index of solvent (<i>n</i>)	QΥ (Φ)
Rhodamine B	27640	0.0043	1.362	0.65

Table S1. QYs of the MoS₂ QDs and aggregated MoS₂ QDs dispersed in ethanol.



Figure S12. ¹H NMR spectra of TPE-CHO in CDCl₃. ¹H NMR (CDCl₃, 400 MHz, ppm): δ 7.02-7.03 (m, 3H), 7.11 (m, 12H), 7.18 (d, 2H), 7.60-7.63 (d, 2H), 9.90 (s, 1H).



Figure S13. The digital photographs of (a) TPE on the micelles surfaces in the faint yellow powder stored at room temperature; (b) the blue fluorescence of TPE on the micelles surfaces under UV light ($\lambda_{ex} = 365$ nm).



Figure S14. (a, b) The TEM images obtained by drying the two aqueous dispersions corresponded to the inset in every picture at the same condition. (c) TEM image of the Cu²⁺ dispersion. (d) TEM image of the aggregated micelles.



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