

**Enzyme and pH dual responsive linear-dendritic block
copolymer micelles based on a phenylalanyl-lysine motif
and peripherally ketal-functionalized dendron as potential
drug carriers**

Yujia Wang,[‡] Wenjie Song,[‡]¹ Lijun Bao, Junwu Wei,

Yangyang Qian, Yunmei Bi*

College of Chemistry and Chemical Engineering, Yunnan Normal University,

Kunming 650092, P.R. China

¹ [‡] These authors contributed equally to this work.

*E-mail: yunmeibi@hotmail.com

Spectroscopic Characterization

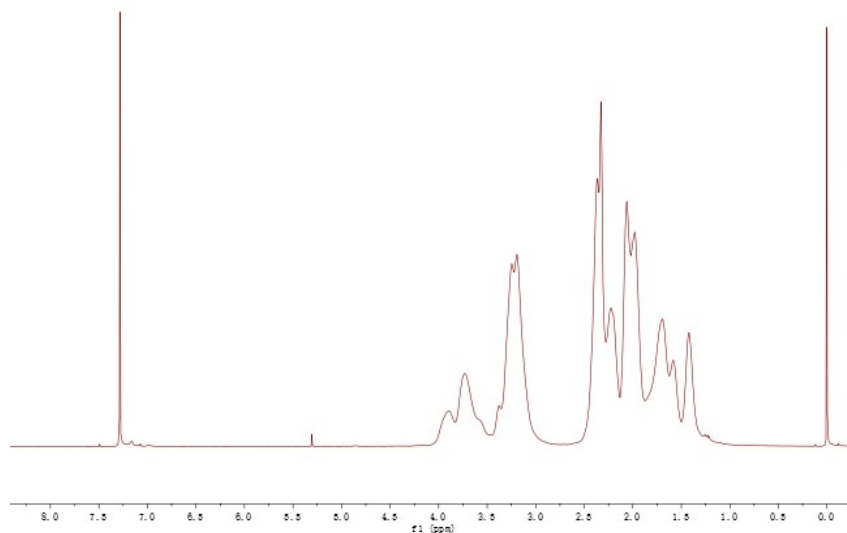


Fig. S1 ¹H NMR spectrum of PNVP-Phe-Lys-*b*-G₁ in CDCl₃.

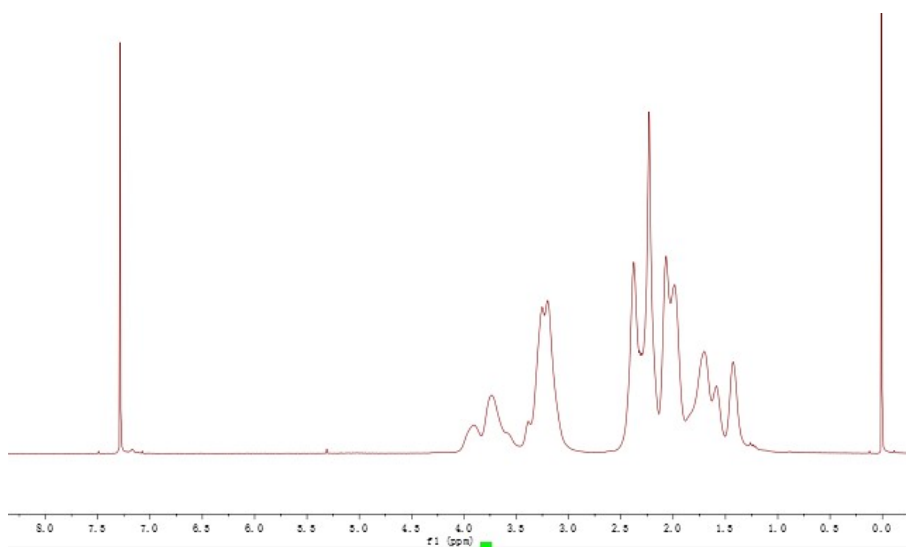


Fig. S2 ¹H NMR spectrum of PNVP-Phe-Lys-*b*-G₂ in CDCl₃.

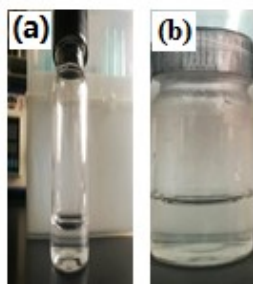


Fig. S3 PNVP-Phe-Lys-*b*-G₃ micelle solutions before and after the incubation with trypsin solution (75 μ M) in 37 $^{\circ}$ C for 48 h.

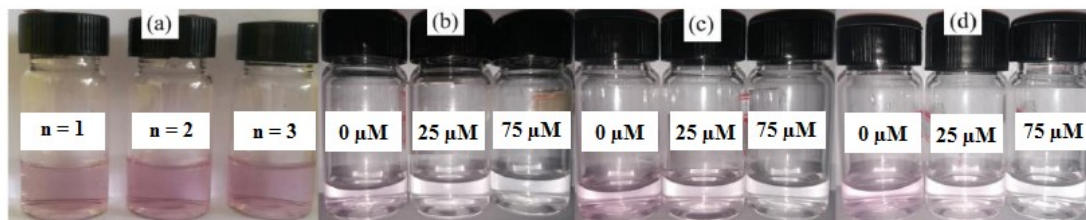


Fig. S4 Nile red loaded PNVP-Phe-Lys-*b*-G_n (n=1-3) micelle solutions (a) and after the incubation with different concentrations of trypsin (0 μ M, 25 μ M, 75 μ M) at 37 $^{\circ}$ C for 48 h ((b) for PNVP-Phe-Lys-*b*-G₁, (c) for PNVP-Phe-Lys-*b*-G₂, (d) for PNVP-Phe-Lys-*b*-G₃).

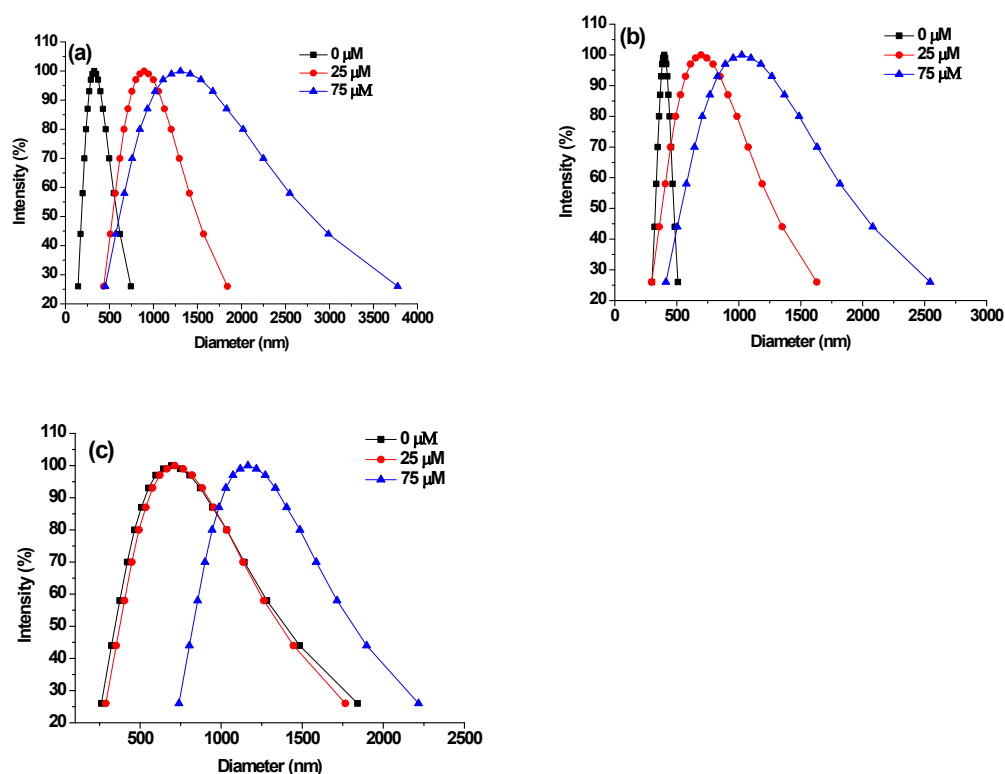


Fig. S5 The particle size of Nile red loaded micelles formed by PNVP-Phe-Lys-*b*-G₁ (a), PNVP-Phe-Lys-*b*-G₂ (b) and PNVP-Phe-Lys-*b*-G₃ (c) after the incubation with different concentrations of trypsin (0 μ M, 25 μ M and 75 μ M).

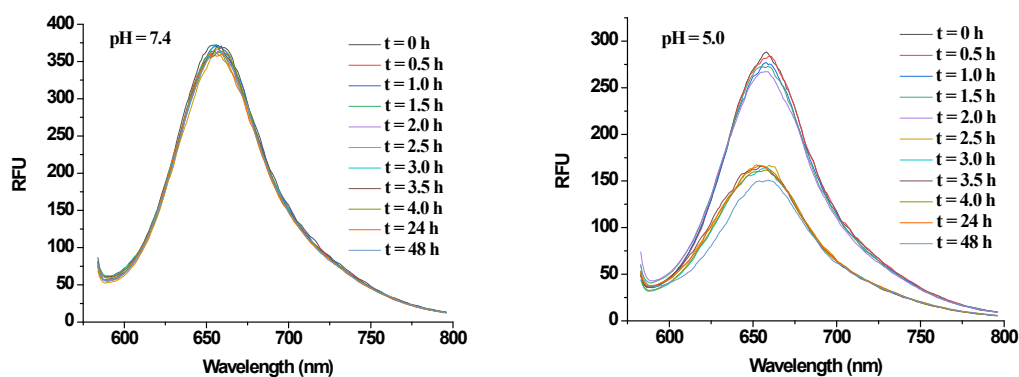


Fig. S6 Fluorescence spectra of Nile red in PNVP-Phe-Lys-*b*-G₁ micelles at pH 7.4 and pH 5.0.

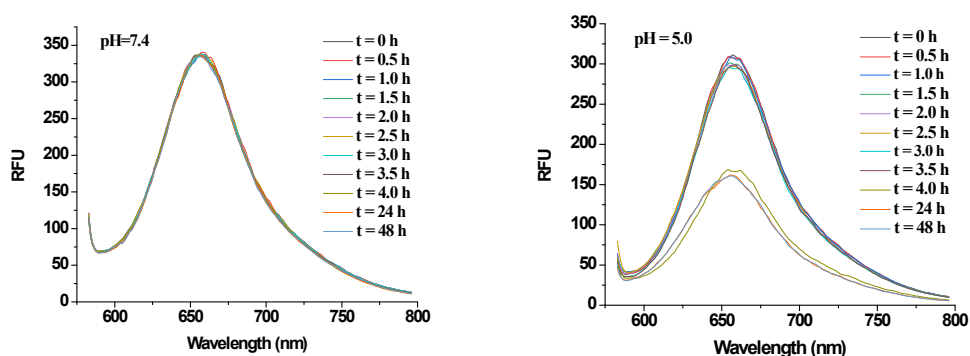


Fig. S7 Fluorescence spectra of Nile red in PNVP-Phe-Lys-*b*-G₂ micelles at pH 7.4 and pH 5.0.

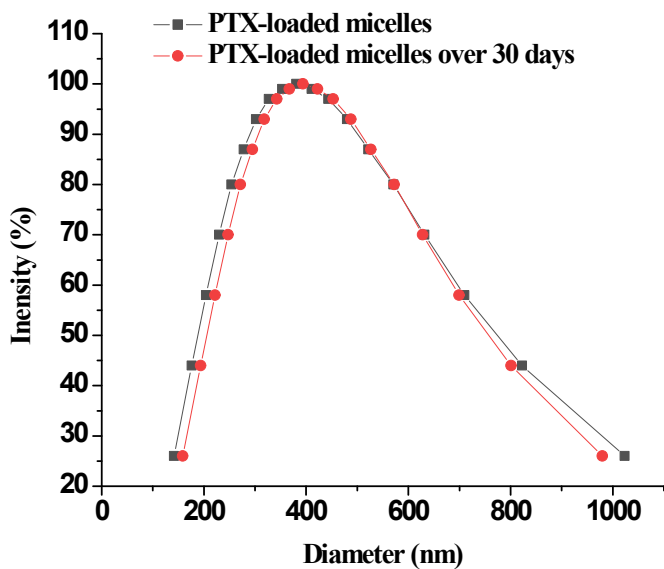


Fig. S8 The particle size of PTX-loaded micelles formed by PNVP-Phe-Lys-*b*-G₃.