

Supporting Information for

**Thermo-Reversible Capture and Release of DNA by
Zwitterionic Surfactants**

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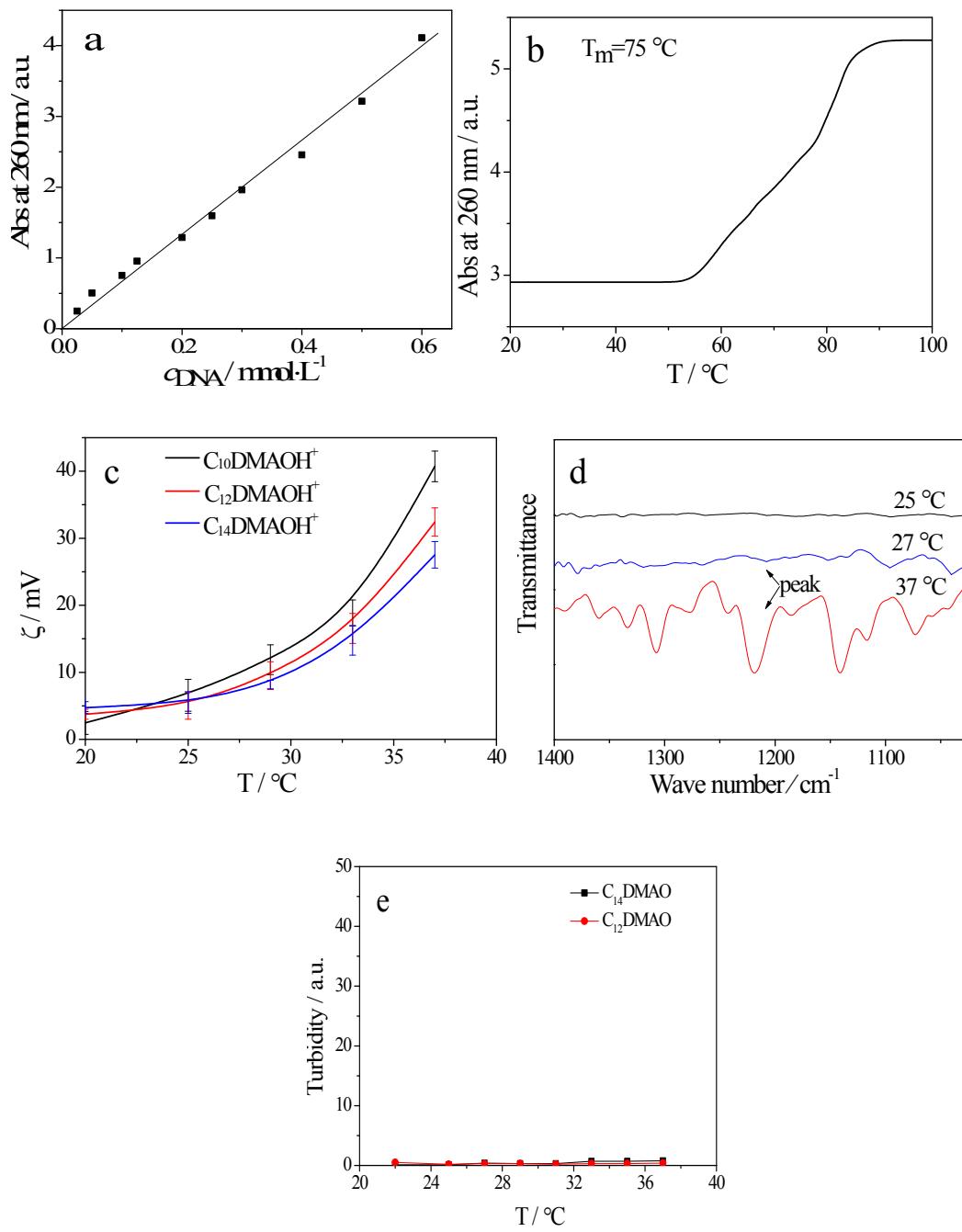


Figure S1. a) Variation of the intensity of the absorbance at 260 nm of DNA as a function in tris-HCl buffer solution with a pH of 6.8 at 25 °C; b) melting curves obtained in the presence of a tris-HCl buffer solution, pH=6.8, $c_{\text{DNA}} = 0.5 \text{ mmol}\cdot\text{L}^{-1}$; c) Variation of zeta-potential (ζ) for $C_n\text{DMAOH}^+$ samples with varying temperatures, samples of $C_{14}\text{DMAO}$, $C_{12}\text{DMAO}$ and $C_{10}\text{DMAO}$ were prepared using tris-HCl buffer solutions with pH = 7.2, 6.8 and 6.6, respectively; d) FT-IR spectra of

C_{14} DAMO in tris-HCl solutions at different temperatures; e) Turbidity of C_{14} DAMO and C_{12} DAMO in tris-HCl solutions at different temperatures.

Table 1. cmcs of C_n DMAO in tris-HCl buffer solutions.

| | ^a cmc | ^b cmc | ^c cmc |
|---------------|---------------------------|--------------------------|--------------------------|
| C_{10} DMAO | 6 mmol·L ⁻¹ | 10 mmol·L ⁻¹ | 15 mmol·L ⁻¹ |
| C_{12} DMAO | 0.9 mmol·L ⁻¹ | 1.6 mmol·L ⁻¹ | 4 mmol·L ⁻¹ |
| C_{14} DMAO | 0.25 mmol·L ⁻¹ | 0.4 mmol·L ⁻¹ | 0.6 mmol·L ⁻¹ |

Samples of C_{14} DMAO, C_{12} DMAO and C_{10} DMAO were prepared using tris-HCl buffer solutions with pH = 7.2, 6.8 and 6.6, respectively.

^a CMC: The onset concentration for the change of I_1/I_3 ratio

^b CMC: The mid-point concentration for the change of I_1/I_3 ratio

^c CMC: The leveling off concentration for the change of I_1/I_3 ratio

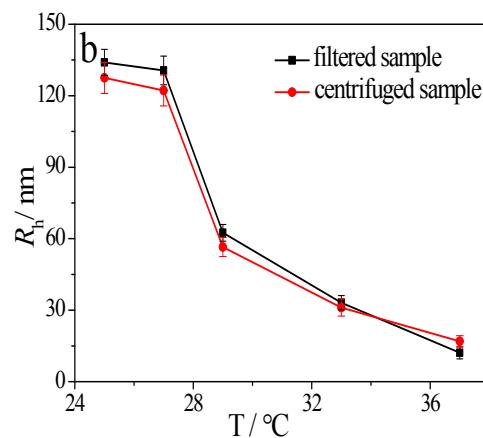


Figure S2. A comparison in hydrodynamic radii (R_h) of DNA/ C_{12} DMAO system at different temperatures for the filtered and centrifuged samples. $c_{DNA} = 0.5$ mmol·L⁻¹ and $R = 20$.

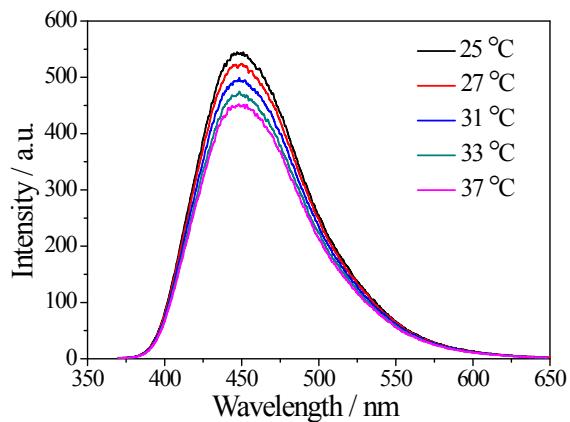


Figure S2. Fluorescence spectra of $0.5 \text{ mmol}\cdot\text{L}^{-1}$ DNA in tris-HCl buffer solution at different temperatures. The samples were prepared using tris-HCl buffer solution with $\text{pH} = 7.2$.

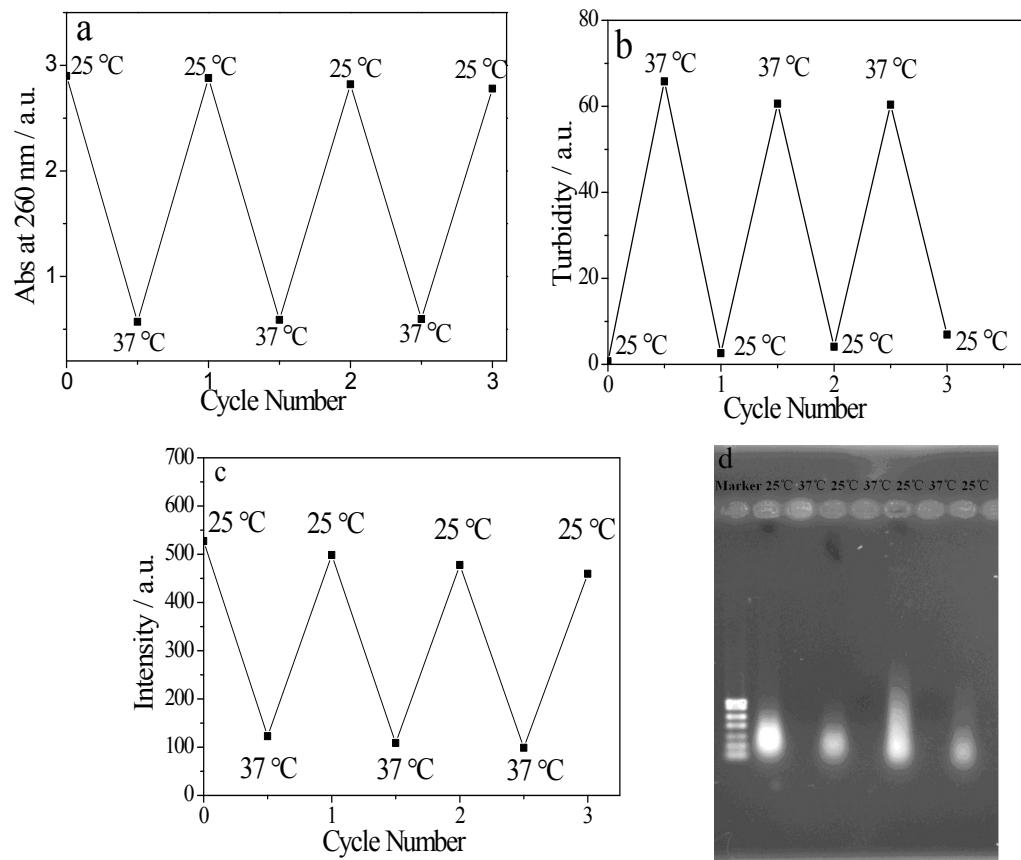


Figure S4. Variations of a) UV-vis absorbance, b) turbidity, c) fluorescence intensity (450 nm) and d) gel electrophoresis performance of DNA/C₁₄DMAO system at 25 °C and 37 °C as a function of cycle number. The samples were prepared using tris-HCl buffer solutions with $\text{pH} = 7.2$.