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Supplementary Information

Amphiphilic Cationic Copolymers with Ciprofloxacin:

Preparation and Antimicrobial Activities

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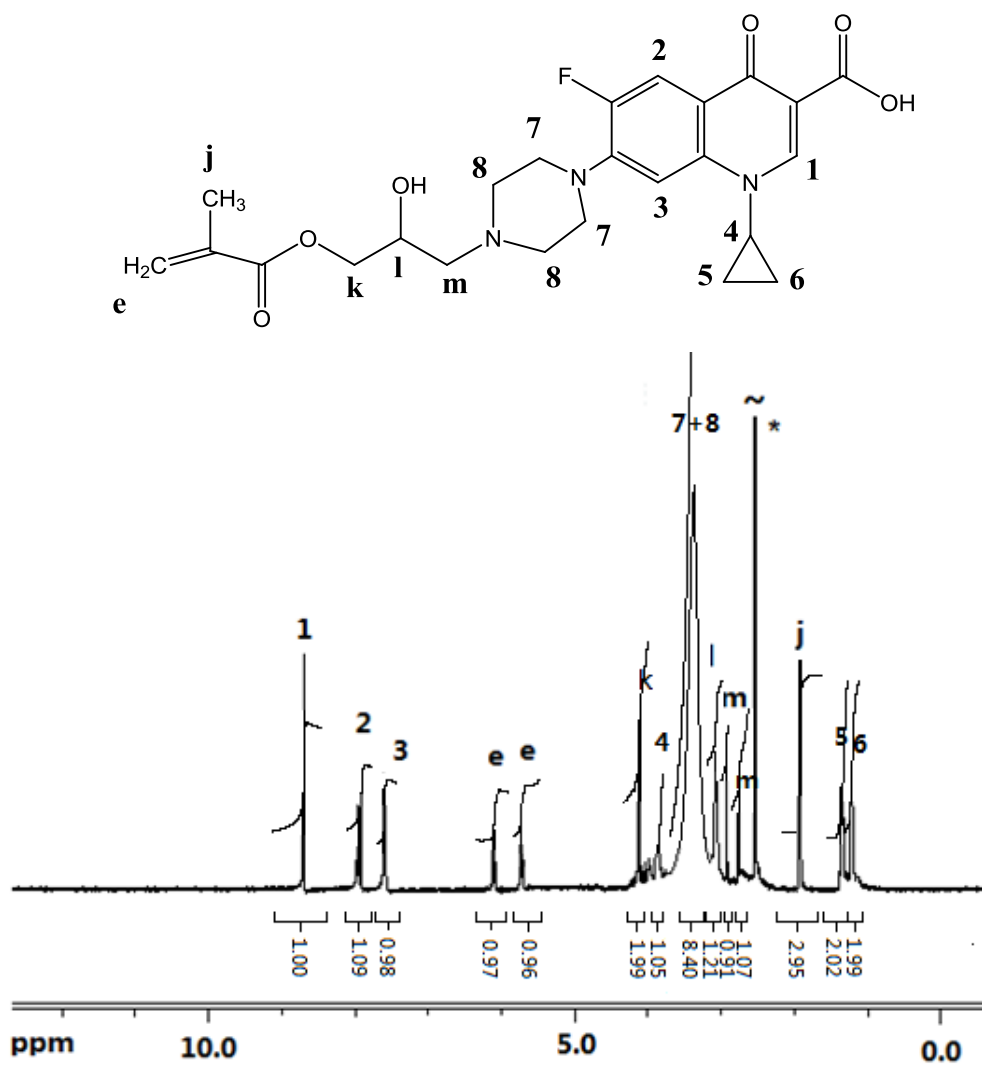


Figure S1. ¹H-NMR spectrum of GMA-CPF in DMSO-d₆

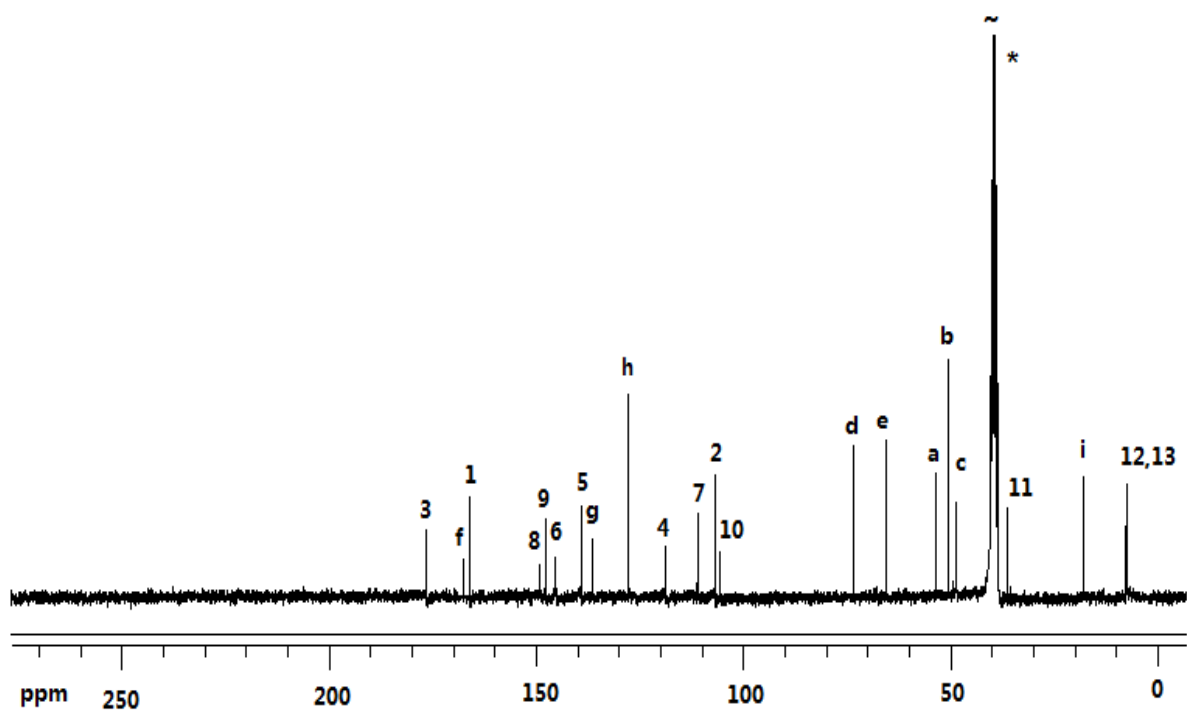
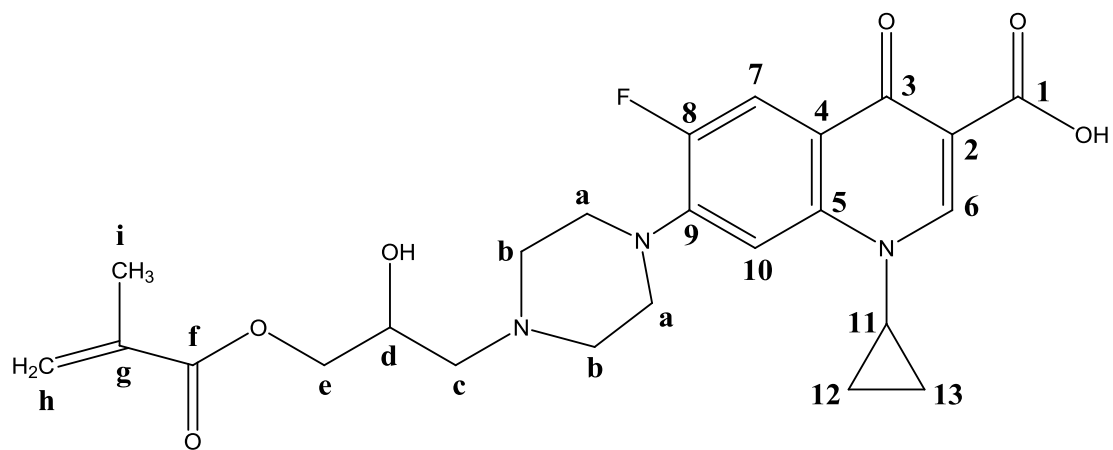


Figure S2. ^{13}C -NMR spectrum of GMA-CPF

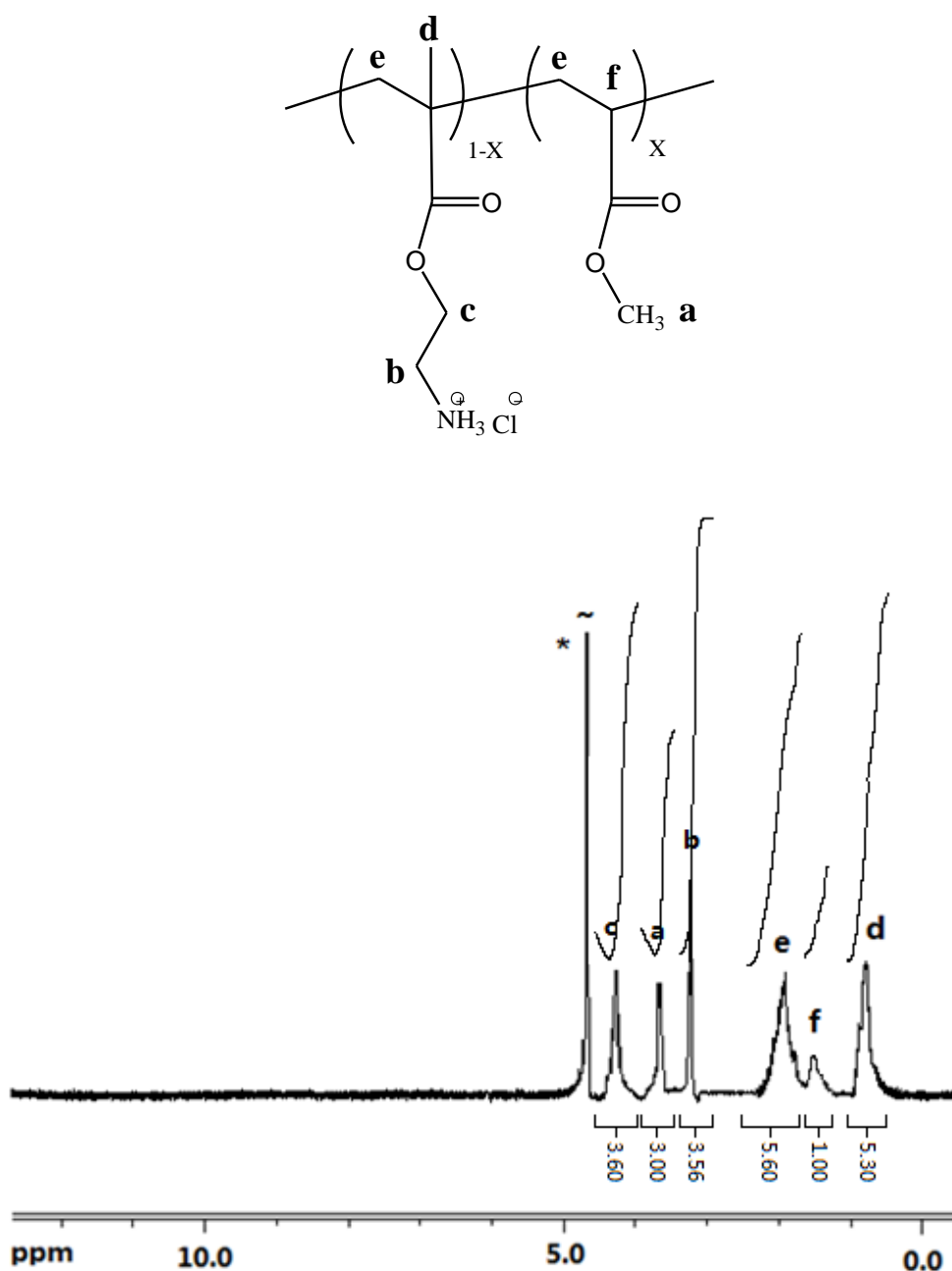


Figure S3. ¹H-NMR spectrum of AEMA/MA copolymer in DMSO-d₆;
(1-x): x=64: 36

The copolymer compositions for this system were determined by ¹H-NMR spectroscopy analysis. The ¹H-NMR peak at 3.65 ppm (Figure S3, a) corresponds to the methyl group of MA side chain; the peak at 3.27 ppm (Figure S3, b) corresponds to the methylene group, -O-CH₂-CH₂-NH₃⁺Cl⁻, in the AEMA side chain. Let m_1 be the mole fraction of AEMA and $m_2 = (1-m_1)$ that of the MA unit.

$$\frac{Area_{3.65ppm}}{Area_{3.27ppm}} = \frac{3m_2}{2m_1} = C_1 \quad \text{Equation 1a}$$

Where $\text{Area}_{3.65\text{ppm}}$ =area of the peak at 3.65ppm and $\text{Area}_{3.27\text{ppm}}$ =area of the peak at 3.27ppm.

On simplification,

$$m_1 = \frac{3}{2C_1+3} \quad \text{Equation 1b}$$

Where m_1 and m_2 are the copolymer molar compositions. The results are presented in Table 1, which are in agreement with the values calculated by apparent charge density.

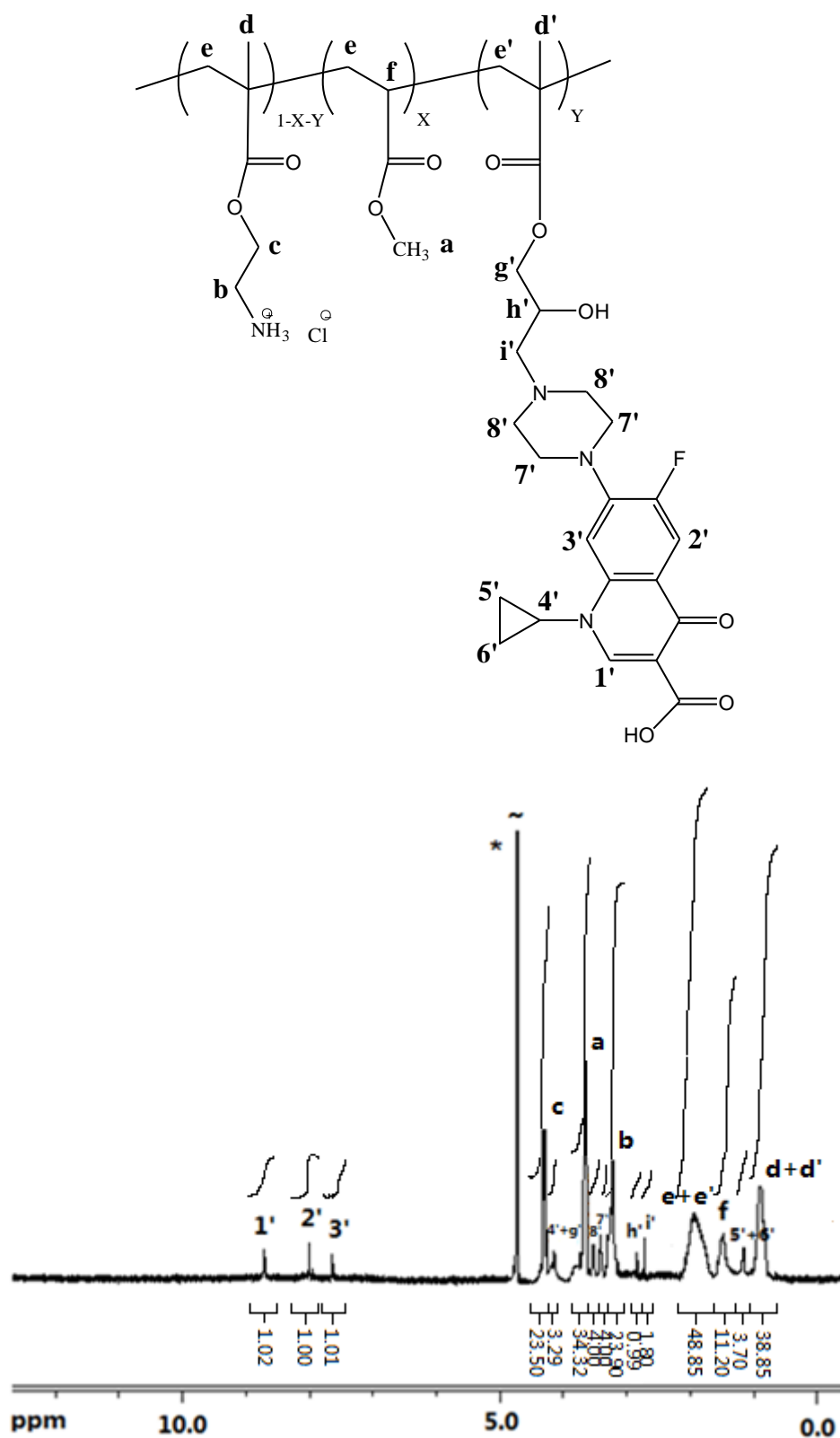


Figure S4. ¹H-NMR spectrum of AEMAH/MA/GMA-CPF copolymer in DMSO-d₆; (1-x-y): x: y=49: 46.9: 4.1

The copolymer compositions for this system were determined by ¹H-NMR

spectroscopy analysis. The $^1\text{H-NMR}$ peak at 3.65 ppm (Figure S4, a) corresponds to the methyl group of MA side chain; the peak at 3.27 ppm (Figure S4, b) corresponds to the methylene group, $-\text{O-CH}_2\text{-CH}_2\text{-NH}_3^+\text{Cl}^-$, in the AEMAH side chain; the peaks at 0.81 ppm (Figure S4, d+d') are the methyl group for AEMAH and GMA-CPF. Considering the mole fraction of AEMAH as m_1 , the mole fraction of MA as m_2 , and $m_3 = (1-m_1-m_2)$ that of the GMA-CPF unit.

$$\frac{\text{Area}_{3.65\text{ppm}}}{\text{Area}_{3.27\text{ppm}}} = \frac{3m_2}{2m_1} = C_1 \quad \text{Equation 2a}$$

$$\frac{\text{Area}_{3.27\text{ppm}}}{\text{Area}_{0.81\text{ppm}}} = \frac{2m_1}{3m_1+3m_3} = C_2 \quad \text{Equation 2b}$$

Where $\text{Area}_{3.65\text{ppm}}$ =area of the peak at 3.65ppm, $\text{Area}_{3.27\text{ppm}}$ =area of the peak at 3.27ppm, and $\text{Area}_{0.81\text{ppm}}$ =area of the peak at 0.81ppm.

On simplification,

$$m_1 = \frac{3C_2}{2C_1C_2+2} \quad \text{Equation 2c}$$

$$m_2 = \frac{C_1C_2}{C_1C_2+1} \quad \text{Equation 2d}$$

$$m_3 = \frac{2-3C_2}{2C_1C_2+2} \quad \text{Equation 2e}$$

Where m_1 , m_2 , and m_3 are the copolymer molar compositions. The results are presented in Table 1, which are in agreement with the values calculated by apparent charge density and UV measurements.