

Supplementary Material

Characterizations of *N*-methylated PEI (1)

FT-IR (cm^{-1}): 2953 and 2833 (C-H str), 1458 (C-H bend), $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 2.25 (s, 3H, $-\text{N}(\text{CH}_3)-$), 2.43-2.6 (m, 4H, $-\text{N}(\text{CH}_2\text{CH}_2)-$).

Characterizations of activated esters and amides intermediates (2a-2c and 3a-3c)

Butyl 2-bromoethanoate (2a): Yield, 97%; $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 0.89 (t, 3H terminal $-\text{CH}_3$), 1.34 (m, 2H, $-\text{CH}_2\text{CH}_2\text{CH}_3$), 1.56 (m, 2H, $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$), 3.88 (s, 2H, $-\text{COCH}_2\text{Br}$), 4.18 (t, 2H, $-\text{COOCH}_2-$).

Hexyl 2-bromoethanoate (2b): Yield, 98%; $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 0.88 (t, 3H, terminal $-\text{CH}_3$), 1.28 (m, 6H, $-\text{CH}_2(\text{CH}_2)_3-$), 1.61 (m, 2H, $-\text{CH}_2(\text{CH}_2)_3\text{CH}_3$), 3.89 (s, $-\text{COCH}_2\text{Br}$, 2H), 4.14 (t, 2H, $-\text{COOCH}_2-$).

Octyl 2-bromoethanoate (2c): Yield, 95%; $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 0.89 (t, 3H, terminal $-\text{CH}_3$), 1.28 (m, 10H $-\text{CH}_2(\text{CH}_2)_5\text{CH}_3$), 1.62 (m, 2H, $-\text{CH}_2(\text{CH}_2)_5\text{CH}_3$), 3.92 (s, 2H, $-\text{COCH}_2\text{Br}$), 4.20 (t, 2H, $-\text{COOCH}_2-$).

***N*-butyl-2-bromoethanamide (3a):** Yield, 100%; $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 0.88 (t, 3H, terminal $-\text{CH}_3$), 1.35 (m, 2H, $-\text{CH}_2\text{CH}_2\text{CH}_3$), 1.56 (m, 2H, $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$), 3.29 (q, 2H, $-\text{CONHCH}_2-$), 3.92 (s, 2H, $-\text{COCH}_2\text{Br}$), 6.72 (br. s, 1H, amide $-\text{NHCO}-$).

***N*-hexyl-2-bromoethanamide (3b):** Yield, 96%; $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 0.91 (t, terminal 3H, $-\text{CH}_3$), 1.3 (m, 6H, $-\text{CH}_2(\text{CH}_2)_3-$), 1.49 (m, 2H, $-\text{CH}_2(\text{CH}_2)_3\text{CH}_3$), 3.31 (q, 2H, $-\text{CONHCH}_2-$), 3.88 (s, 2H, $-\text{COCH}_2\text{Br}$), 6.54 (br. s, 1H, amide $-\text{NHCO}-$).

***N*-octyl-2-bromoethanamide (3c) :** Yield, 95.5%; $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ/ppm 0.88 (t, terminal 3H, $-\text{CH}_3$), 1.28 (m, 10H $-\text{CH}_2(\text{CH}_2)_5\text{CH}_3$), 1.61 (m, 2H, $-\text{CH}_2(\text{CH}_2)_5\text{CH}_3$), 3.29 (q, 2H, $-\text{CONHCH}_2-$), 3.94 (s, 2H, $-\text{COCH}_2\text{Br}$), 7.22 (br. s, 1H, amide $-\text{NHCO}-$).

Characterizations of the ACM- N_{alk} (4-7)

ACM-N_{But} (4): Yield, 83.4%; ¹H NMR (400 MHz, D₂O): δ/ppm 0.96 (t, 3H, terminal -CH₃), 1.38-1.51 (br, 2H, -CH₂CH₂CH₃), 1.72-1.89 (br, 2H, -CH₂CH₂CH₂N(CH₃)CH₂CH₂-), 3.12-3.6 (br, 9H, -(CH₂)₂N(CH₂)(CH₃)-).

ACM-N_{Hex} (5): Yield, 84%; ¹H NMR (400 MHz, D₂O): δ/ppm 0.85 (t, 3H, terminal -CH₃), 1.18-1.48 (br, 6H, -CH₂(CH₂)₃-), 1.70-1.85 (br, 2H, -CH₂CH₂N(CH₃)CH₂CH₂-), 3.41-4.09 (br, 9H, -(CH₂)₂N(CH₂)(CH₃)-).

ACM-N_{Oct} (6): Yield, 82.5%; ¹H NMR (400 MHz, D₂O): δ/ppm 0.88 (t, 3H, terminal -CH₃), 1.16-1.50 (br, 10H, -CH₂(CH₂)₅-), 1.59-1.89 (br, 2H, -CH₂CH₂N(CH₃)CH₂CH₂-), 3.17-4.44 (br, 9H, -(CH₂)₂N(CH₂)(CH₃)-).

ACM-N_{Dec} (7): Yield, 80.3%; ¹H NMR (400 MHz, CDCl₃): δ/ppm 0.88 (t, 3H, terminal -CH₃), 1.22-1.48 (br, 14H, -CH₂(CH₂)₇-), 1.6-1.75 (br, 2H, -CH₂CH₂N(CH₃)CH₂CH₂-), 3.3-4.04 (br, 9H, -(CH₂)₂N(CH₂)(CH₃)-).

Characterizations of the ACM-E_{alk} (8-10) and ACM-A_{alk} (11-13)

ACM-E_{But} (8): Yield, 76.5%; ¹H NMR (400 MHz, D₂O): δ/ppm 0.88 (t, 3H, terminal -CH₃), 1.22-1.35 (br, 2H, -CH₂CH₂CH₃), 1.65-1.75 (q, 2H, -CH₂CH₂CH₂N(CH₃)CH₂CH₂-), 3.52-3.85 (br, 7H, -(CH₂)₂N(CH₂)(CH₃)-), 3.9-4.4 (br, 2H, -CH₂OCOCH₂N(CH₃)CH₂CH₂-), 4.5-4.75 (br, 2H, -OCOCH₂N(CH₃)CH₂CH₂-).

ACM-E_{Hex} (9): Yield, 76%; ¹H-NMR (400 MHz, CDCl₃): δ/ppm 0.89 (t, 3H, terminal -CH₃), 1.24-1.39 (br m, 6H, -CH₂(CH₂)₃-), 1.7-1.85 (q, 2H, -CH₂CH₂N(CH₃)CH₂CH₂-), 3.79-4.29 (m, 7H, -(CH₂)₂N(CH₂)(CH₃)-), 4.35-4.6 (m, 2H, -CH₂OCOCH₂N(CH₃)CH₂CH₂-), 4.55-4.74 (m, 2H, -OCOCH₂N(CH₃)CH₂CH₂-).

ACM-E_{Oct} (10): Yield, 75.4%; ¹H NMR (400 MHz, CDCl₃): δ/ppm 0.87 (t, 3H, terminal -CH₃), 1.28-1.38 (br, 10H, -CH₂(CH₂)₅-), 1.75-1.87 (q, 2H, -CH₂CH₂N(CH₃)CH₂CH₂-), 3.6-3.8 (br, 7H, -(CH₂)₂N(CH₂)(CH₃)-), 4.1-4.45 (br, 2H, -CH₂OCOCH₂N(CH₃)CH₂CH₂-), 4.55-4.89 (br, 2H, -OCOCH₂N(CH₃)CH₂CH₂-).

ACM-A_{But} (11): Yield, 76.5%; ¹H NMR (400 MHz, D₂O): δ/ppm 0.85 (t, 3H, terminal -CH₃), 1.28-1.48 (br, 2H, -CH₂CH₂CH₃), 1.82 (q, 2H, -CH₂CH₂CH₂N(CH₃)CH₂CH₂-), 3.1-3.8 (br, 7H, -

$(CH_2)_2N(CH_2)(CH_3)-$, 3.9-4.4 (br, 2H, $-CH_2NHCOCH_2N(CH_3)CH_2CH_2-$), 4.5-4.9 (br, 2H, $-NHCOCH_2N(CH_3)CH_2CH_2-$), 8.0-8.5 (br, 1H, $-CONHCH_2-$).

ACM-A_{Hex} (12): Yield, 79%; 1H NMR (400 MHz, $CDCl_3$): δ /ppm 0.88 (t, 3H, terminal $-CH_3$), 1.2-1.39 (br, 6H, $-CH_2(CH_2)_3-$), 1.72- 1.9 (q, 2H, $-CH_2CH_2CH_2N(CH_3)CH_2CH_2-$), 3.2-3.89 (br, 7H, $(CH_2)_2N(CH_2)(CH_3)-$), 4.05-4.3 (br, 2H, $-CH_2NHCOCH_2N(CH_3)CH_2CH_2-$), 4.5-4.86 (br, 2H, $-NHCOCH_2N(CH_3)CH_2CH_2-$), 8.05-8.5 (br, 1H, $-CONHCH_2-$).

ACM-A_{Oct} (13): Yield, 75.2%; 1H NMR (400 MHz, $CDCl_3$): δ /ppm 0.89 (t, 3H, terminal $-CH_3$), 1.2-1.49 (br m, 10H, $-CH_2(CH_2)_5$), 1.7-1.87 (q, 2H, $-CH_2CH_2CH_2N(CH_3)CH_2CH_2-$), 3.1-3.67 (br, 7H, $(CH_2)_2N(CH_2)(CH_3)-$), 3.93-4.3 (br, 2H, $-CH_2NHCOCH_2N(CH_3)CH_2CH_2-$), 4.6-4.8 (br, 2H, $-NHCOCH_2N(CH_3)CH_2CH_2-$), 8-8.5 (br, 1H, $-CONHCH_2-$).

Table S1. Molecular weight of the amphiphilic cationic macromolecules (ACMs).

Macromolecules	^aDegree of quaternization (DQ) (%)	^bMolecular Weight (M_n) (kDa)
ACM-N_{but} (4)	100	3.6
ACM-N_{Hex} (5)	100	4.2
ACM-N_{Oct} (6)	100	4.7
ACM-N_{Dec} (7)	100	5.2
ACM-E_{But} (8)	100	4.7
ACM-E_{Hex} (9)	100	5.2
ACM-E_{Oct} (10)	100	5.7
ACM-A_{But} (11)	100	4.7
ACM-A_{Hex} (12)	100	5.2
ACM-A_{Oct} (13)	100	5.7

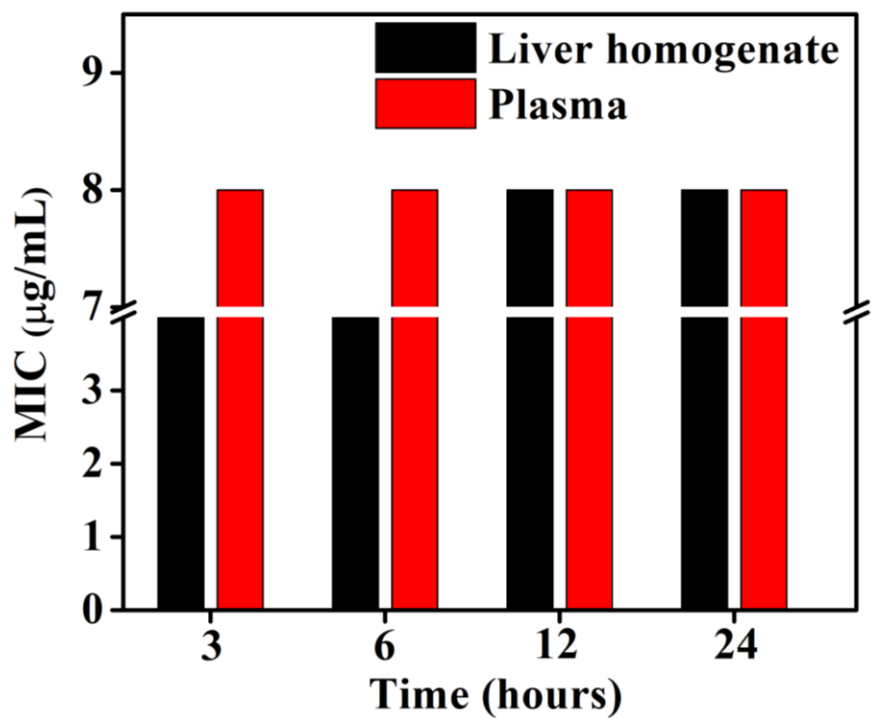


Figure S1. Antibacterial activity of ACM-A_{Hex} (**12**) in physiological fluids. Each concentration had triplicate values and the entire experiment was performed twice. The average data was reported.