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Supporting information

Synthesis of cationic *N*-acylated thiazolidine for selective activity against Gram-positive bacteria, and evaluation of *N*-acylation's role in membranedisrupting activity

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Fig. S1A: ¹H NMR of 2-(pyridine-2-yl)thiazolidine



Fig. S1B: ¹³C NMR of 2-(pyridine-2-yl)thiazolidine



Fig. S1C: HRMS of 2-(pyridine-2-yl)thiazolidine



Fig. S2A: ¹H NMR of PyN10Th



Fig. S2B: ¹³C NMR of PyN10Th



Fig. S2C: HRMS of PyN10Th



Fig. S3A: ¹H NMR of PyN12Th



Fig. S3B: ¹³C NMR of PyN12Th



Fig. S3C: HRMS of PyN12Th



Fig. S4A: ¹H NMR of PyN14Th



Fig. S4B: ¹³C NMR of PyN14Th



Fig. S4C: HRMS of PyN14Th



Fig. S5A: ¹H NMR of PyN16Th



Fig. S5B: ¹³C NMR of PyN16Th



Fig. S5C: HRMS of PyN16Th



Fig. S6A: ¹H NMR of PyN18Th



Fig. S6B: ¹³C NMR of PyN18Th



Fig. S6C: HRMS of PyN18Th



Fig. S7A: ¹H NMR of QPyN10Th



Fig. S7B: ¹³C NMR of QPyN10Th



Fig. S7C: HRMS of QPyN10Th



Fig. S8A: ¹H NMR of QPyN12Th



Fig. S8B: ¹³C NMR of QPyN12Th



Fig. S8C: HRMS of QPyN12Th



Fig. S9A: ¹H NMR of QPyN14Th



Fig. S9B: ¹³C NMR of QPyN14Th



Fig. S9C: HRMS of QPyN14Th



Fig. S10A: ¹H NMR of QPyN16Th



Fig. S10B: ¹³C NMR of QPyN16Th



Fig. S10C: HRMS of QPyN16Th



Fig. S11A: ¹H NMR of QPyN18Th



Fig. S11B: ¹³C NMR of QPyN18Th



Fig. S11C: HRMS of QPyN18Th



Fig. S12: Membrane integrity assay at 1X MIC. Effects of QPyNATh on membrane permeability of (A) UPEC and (B) MRSA. Effects of QPyNATh on membrane depolarization of (C) UPEC and (D) MRSA.



Fig. S13: ROS assay at 1X MIC. Effects of QPyNATh on ROS generation in (A) UPEC and (B) MRSA.