

Electronic Supplementary Information (ESI)

Design, synthesis and biocidal effect of novel amine *N*-halamine microspheres based on 2,2,6,6-tetramethyl-4-piperidinol as promising antibacterial agents

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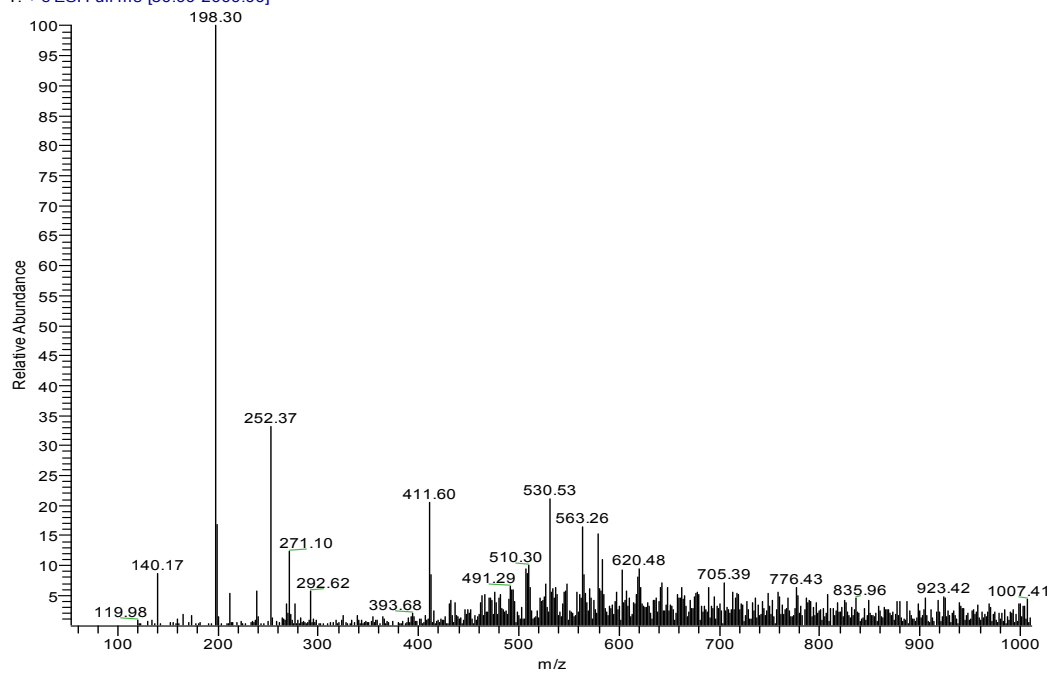


Fig. S1. Mass spectrogram of ATMP

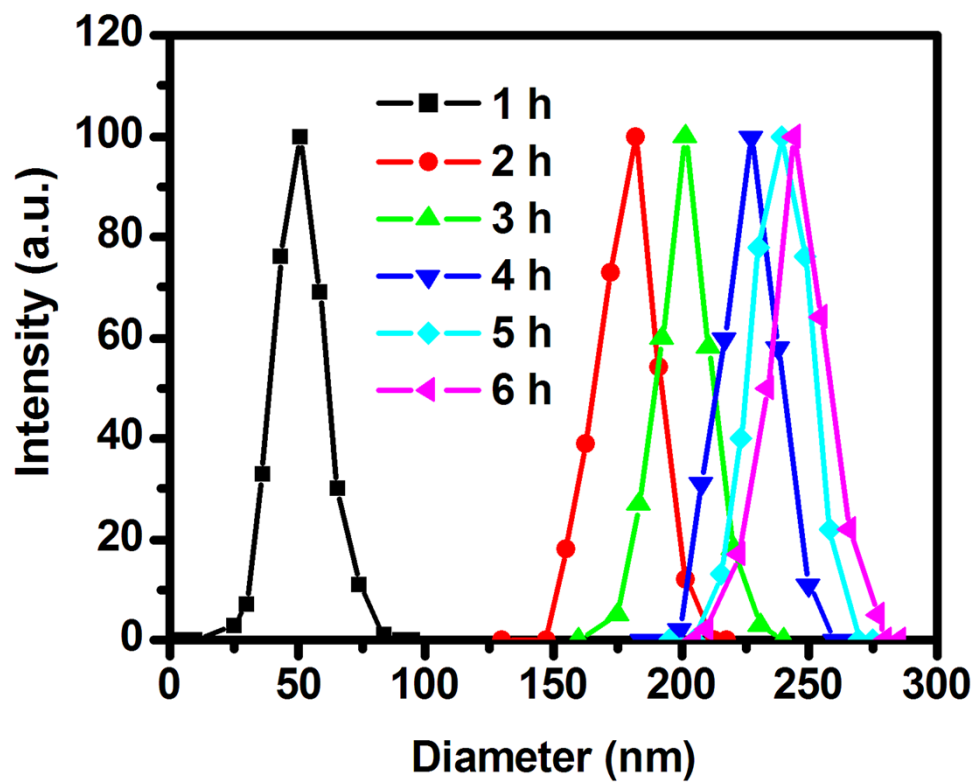
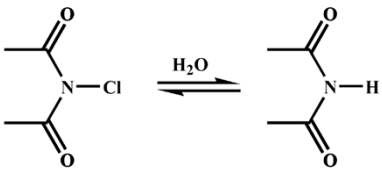
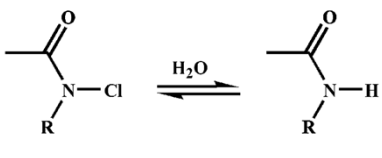
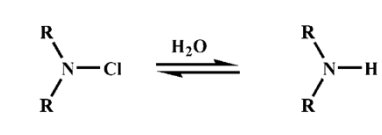


Fig. S2. Particle size distribution of poly(ATMP-co-MMA) microspheres with different copolymerization period.

Table S1. Dissociation constant of different *N*-halamine in aqueous solution

<i>N</i> -halamine type	Dissociation reaction	Dissociation constant ^a
Imide		$< 10^{-4}$
Amide		$< 10^{-9}$
Amine		$< 10^{-12}$

^aDissociation constant was from references.¹⁻⁴

Table S2. Particle size and surface area characteristics of poly(ATMP-co-MMA) microspheres formed with different copolymerization period

Sample	Copolymerization period (h)	Particle size (nm) ^a		Surface area ^b (m ² ·g ⁻¹)
		Size distribution	Average size	
S1	1	20-90	50.7	118.3
S2	2	150-210	181.8	33.0
S3	3	170-240	201.9	29.7
S4	4	190-250	227.5	26.4
S5	5	210-260	239.2	25.1
S6	6	210-280	244.5	24.5

^aParticle size was determined by TEM images.

^bThe surface area was calculated based on the assumption that the particles are non-porous spheres with density of 1.0 g·cm⁻³. The calculation was performed according to the following equation: $S = 6(D \cdot d)^{-1}$, wherein S is the surface area (m²·g⁻¹); D is the diameter (μm); and d is the density (g·cm⁻³) of the particles.⁵

Table S3. Minimum inhibitory concentration (MIC) of different products against *S. aureus*, *B. subtilis*, *E. coli*, and *P. aeruginosa*

Sample	MIC (mg/mL)				Reference
	Gram-positive bacteria		Gram-negative bacteria		
	<i>S. aureus</i>	<i>B. subtilis</i>	<i>E. coli</i>	<i>P. aeruginosa</i>	
PSA@Fe ₃ O ₄ @SiO ₂ - <i>N</i> -halamine ^a	80	-	-	60	6
HMNH NPs ^b	160	160	160	80	7
BAMNH NPs ^c	80	80	80	40	7
H-NHFS NPs ^d	160	-	-	80	8
BA-NHFS NPs ^e	40	-	-	40	8
Amine <i>N</i> -halamine microspheres	10	-	10	-	This study

^aAmide *N*-halamine-immobilized PSA@Fe₃O₄@SiO₂ nanoparticles.

^bHydantoin-structural magnetic amide *N*-halamine nanoparticles.

^cBarbituric acid-based magnetic imide *N*-halamine nanoparticles.

^dHydantoin-originated amide *N*-halamine-functionalized silica nanoparticles.

^eBarbituric acid-originated imide *N*-halamine-functionalized silica nanoparticles.

Table S4. Oxidative chlorine content of amine *N*-halamine microspheres with different soaking period

Soaking period (h)^a	Oxidative chlorine (%)^b	Reduction (%)^c
0	1.17	0
6	1.16	0.85
12	1.16	0.85
18	1.14	2.56
24	1.11	5.13
30	1.10	5.98
36	1.08	7.69
42	1.08	7.69
48	1.07	8.55

^aPeriod since initial soaking.

^bOxidative chlorine content was determined by the iodometric/thiosulfate titration after a certain soaking age.

^cReduction percentage of oxidative chlorine after a certain soaking age.

Supplementary References

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