

Supporting information:

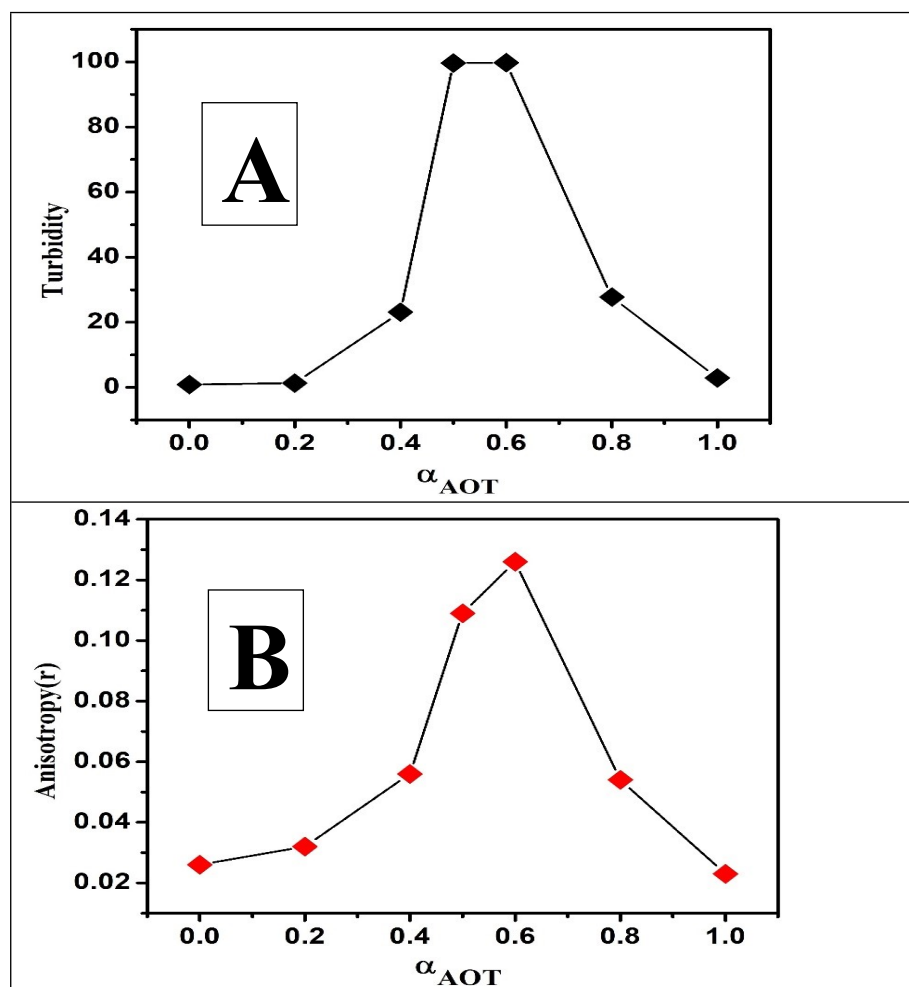
Electrostatic control of Protein-Vesicle interactions in catanionic long-Chain ionic liquid systems: Insights from Human Serum Albumin

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1. Characterization of catanionic vesicle:



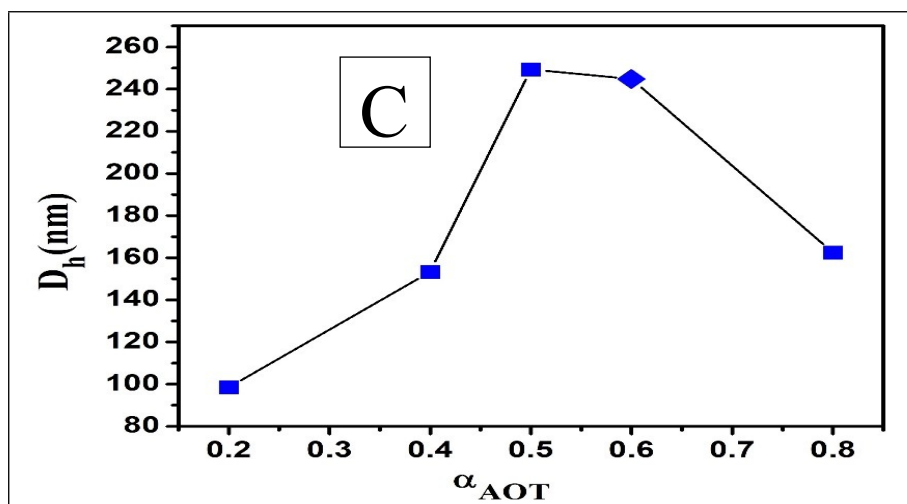


Fig. S1 (A) Variation in turbidity of aqueous [C₁₆Mim] [AOT] mixture at various mole fractions. (B) Variation in steady-state anisotropy of [C₁₆Mim] [AOT] mixture with varying mole fraction of AOT. (C) Change in hydrodynamic diameter (D_h) of [C₁₆Mim] [AOT] aggregate in different mole fractions.

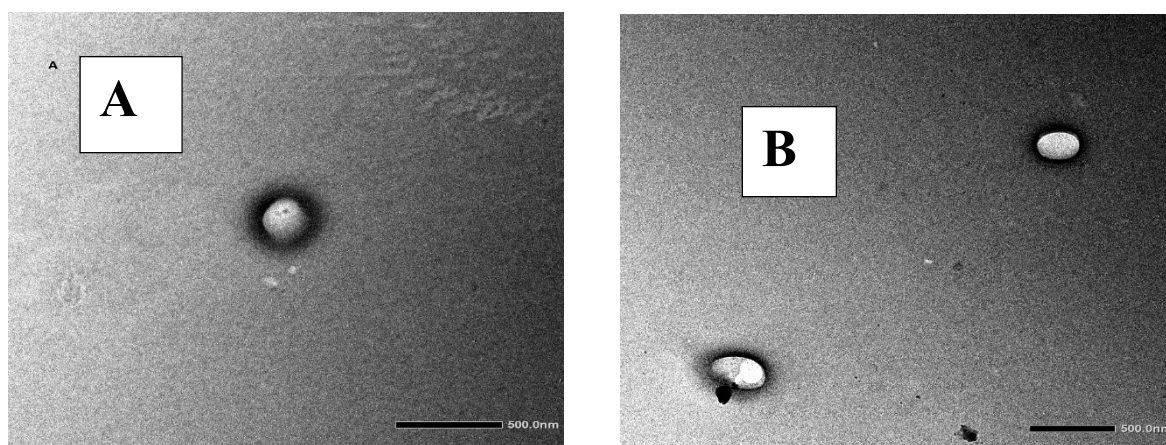


Fig. S2 TEM micrographs of [C₁₆Mim]/[AOT] mixture at (A) 0.5 mole fraction, and (B) 0.6 mole fraction of AOT.

(2) Absorbance spectra of HSA in the presence of [C₁₆Mim] [AOT].

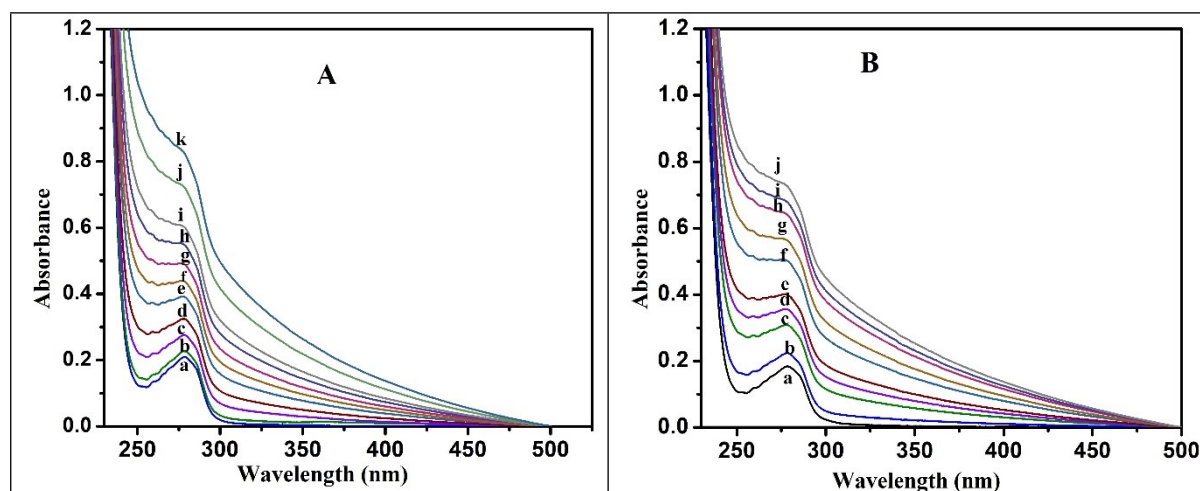


Fig. S3 Absorbance spectra of HSA in the presence of [C₁₆Mim] [AOT], (A) 0.5-mole fraction (a, b, c, d, e, f, g, h, i, j, k; 0.0, 0.05, 0.08, 0.12, 0.15, 0.18, 0.22, 0.28, 0.30, 0.33, 0.35 mM respectively), and (B) 0.6-mole fraction (a, b, c, d, e, f, g, h, i, j; 0.0, 0.04, 0.09, 0.11, 0.13, 0.18, 0.21, 0.26, 0.28, 0.32 mM respectively) of AOT.

(3) Table S1. Time-dependent fluorescence decay profile of HSA in [C₁₆Mim] [AOT] mixed vesicle system for $\chi_{\text{AOT}}=0.5$ and $\chi_{\text{AOT}}=0.6$, respectively.

[C ₁₆ Mim]/[AOT] (mM)($\chi_{\text{AOT}}=0.5$)	τ_1 (ns)	A ₁ (%)	τ_2 (ns)	A ₂ (%)	$\langle \tau \rangle$ (ns)
0	1.64	24.17	5.94	75.83	3.64
0.04	1.39	21.97	5.10	78.03	3.21
0.07	1.03	21.75	5.04	78.25	2.74
0.14	0.61	19.72	4.77	80.82	2.04
0.20	0.21	24.22	4.57	75.78	0.77
0.36	0.15	29.72	4.52	70.28	0.48
0.49	0.071	41.08	4.37	58.92	0.17
[C ₁₆ Mim]/[AOT] (mM)($\chi_{\text{AOT}}=0.6$)					
0.014	1.35	21.86	5.45	78.14	3.28
0.03	1.10	19.23	5.11	80.77	3.04

0.07	1.30	26.63	5.05	73.37	2.86
0.09	1.42	30.21	5.0	69.79	2.85
0.11	1.15	36.12	4.84	63.88	2.24
0.24	0.036	51.82	3.74	48.18	0.07
0.49	0.11	55.49	3.45	44.51	0.19

(4) ITC profile [C₁₆Mim] [AOT] in the presence of HSA solution.

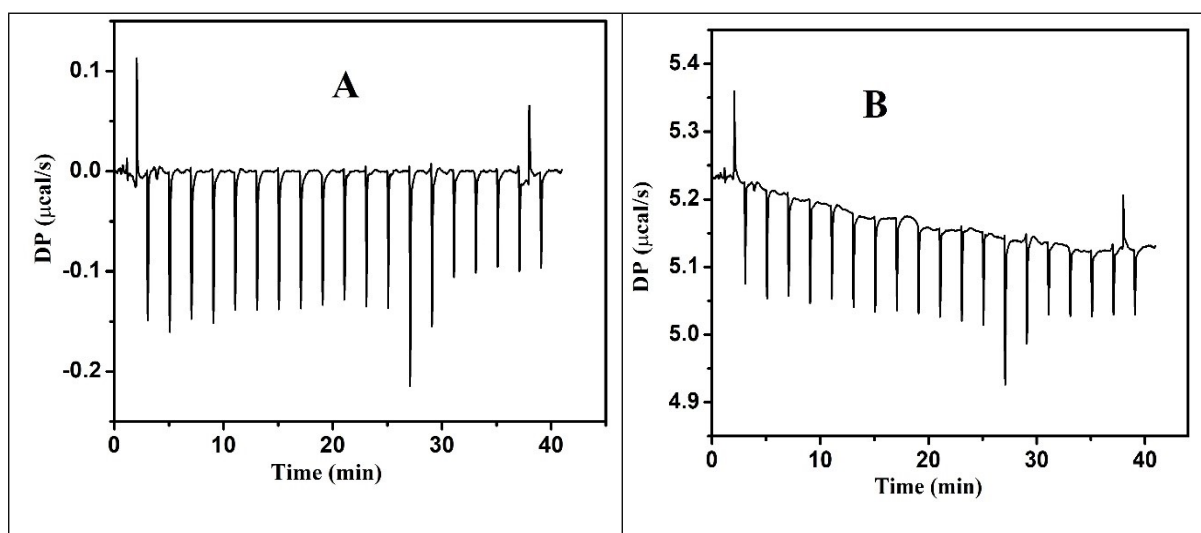


Fig. S4 ITC profile of titration of (A) 0.5, and (B) 0.6- mole fraction of [C₁₆Mim] [AOT] in HSA solution at 298 K.

(5) Variation of hydrodynamic diameter (D_h) of vesicle in the presence of different concentrations of HSA.

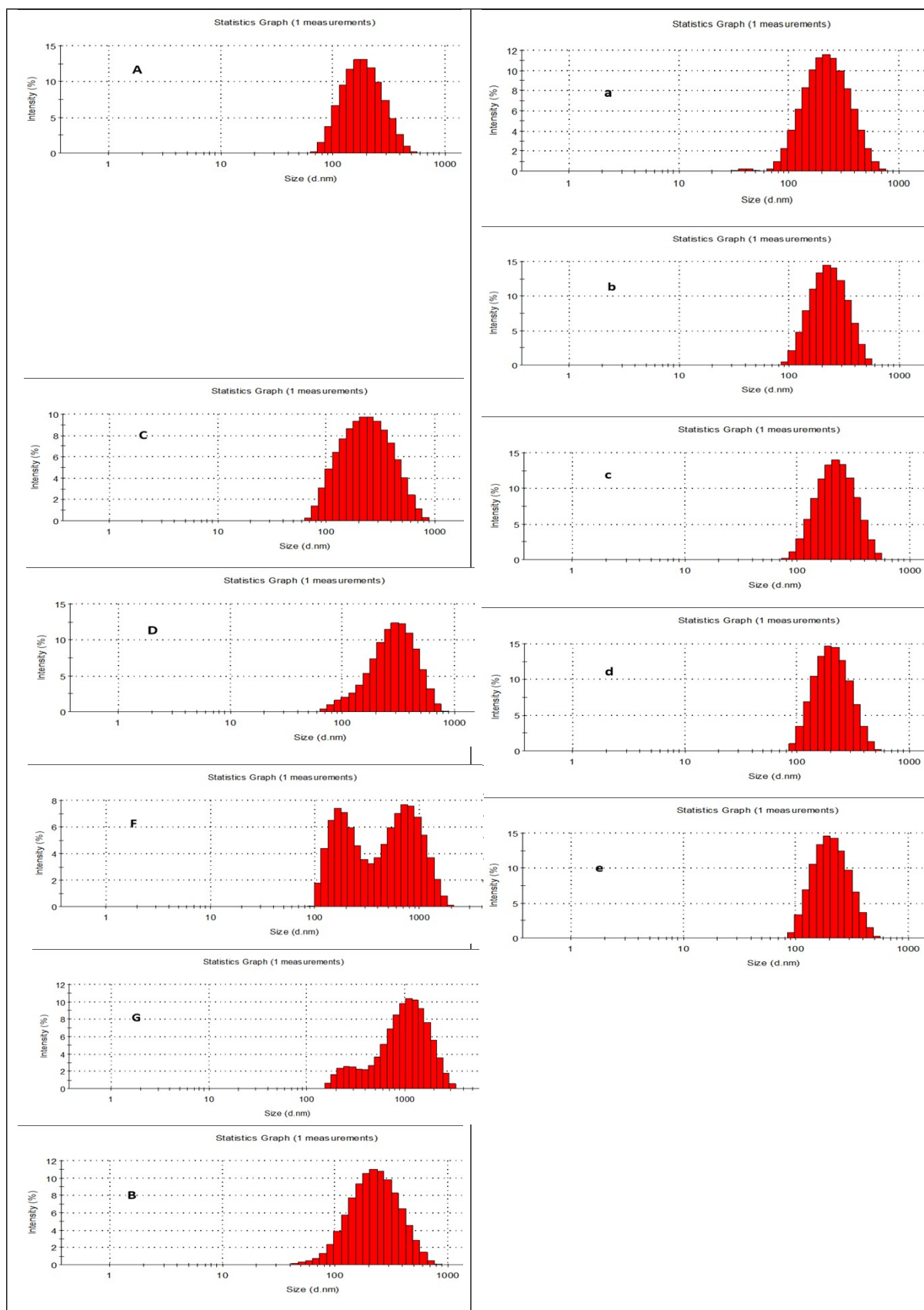


Fig. S5 Variation of hydrodynamic diameter (D_h) of $[C_{16}Mim][AOT]$ vesicle with increasing concentration of HSA; (A, B, C, D, E, F, G; 0.0, 0.36, 0.54, 1.07, 1.42, 2.21, 3.70

μM of HSA for the 0.5-mole fraction of AOT), and (a, b, c, d, e; 0.0, 0.48, 1.13, 2.45, 3.18 μM of HSA for the 0.6-mole fraction of AOT) respectively.

(6) Table S2. Variation of α -helix contents of HSA in the presence of various amounts of $[\text{C}_{16}\text{Mim}] [\text{AOT}]$ (for $\chi_{\text{AOT}}=0.5$ and $\chi_{\text{AOT}}=0.6$) in pH 7.4, at 298 K.

$[\text{C}_{16}\text{Mim}] [\text{AOT}]$ ($\chi_{\text{AOT}}=0.5$) (mM)	α -helix (%)	$[\text{C}_{16}\text{Mim}] [\text{AOT}]$ ($\chi_{\text{AOT}}=0.6$) (mM)	α -helix (%)
0	60.85	0	60.85
0.002	60.85	0.003	59.28
0.011	60.75	0.021	60.21
0.022	60.65	0.052	60.038
0.102	60.53	0.076	59.98
0.107	60.71	0.133	60.09
0.117	60.69	0.136	59.78
0.133	60.48	0.166	59.48
0.186	60.25		