

Electronic supporting information

Complexation of Triblock Reverse Copolymer 10R5 with Surface Active Ionic Liquids in Aqueous Medium: A Physico-chemical Study

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Annexure S1.

The maximum surface excess concentration ($\Gamma_{\max}$) and minimum area per molecule [1, 2] at the air-solution interface can be calculated using equation (1) and (2) as follows:

$$\frac{d\gamma}{d \ln C_T} = -nRT\Gamma_{\max} \quad (1)$$

$$A_{\min} = 10^{20} / (N_A \cdot \Gamma_{\max}) \quad (2)$$

The Gibbs free energy of adsorption ($\Delta G^\circ_{\text{ads}}$) [3] is calculated using the equation (3)

$$\Delta G^\circ_{\text{ads}} = \Delta G_{\text{mic}} - \Pi_{\text{cmc}} / \Gamma_{\max} \quad (3)$$

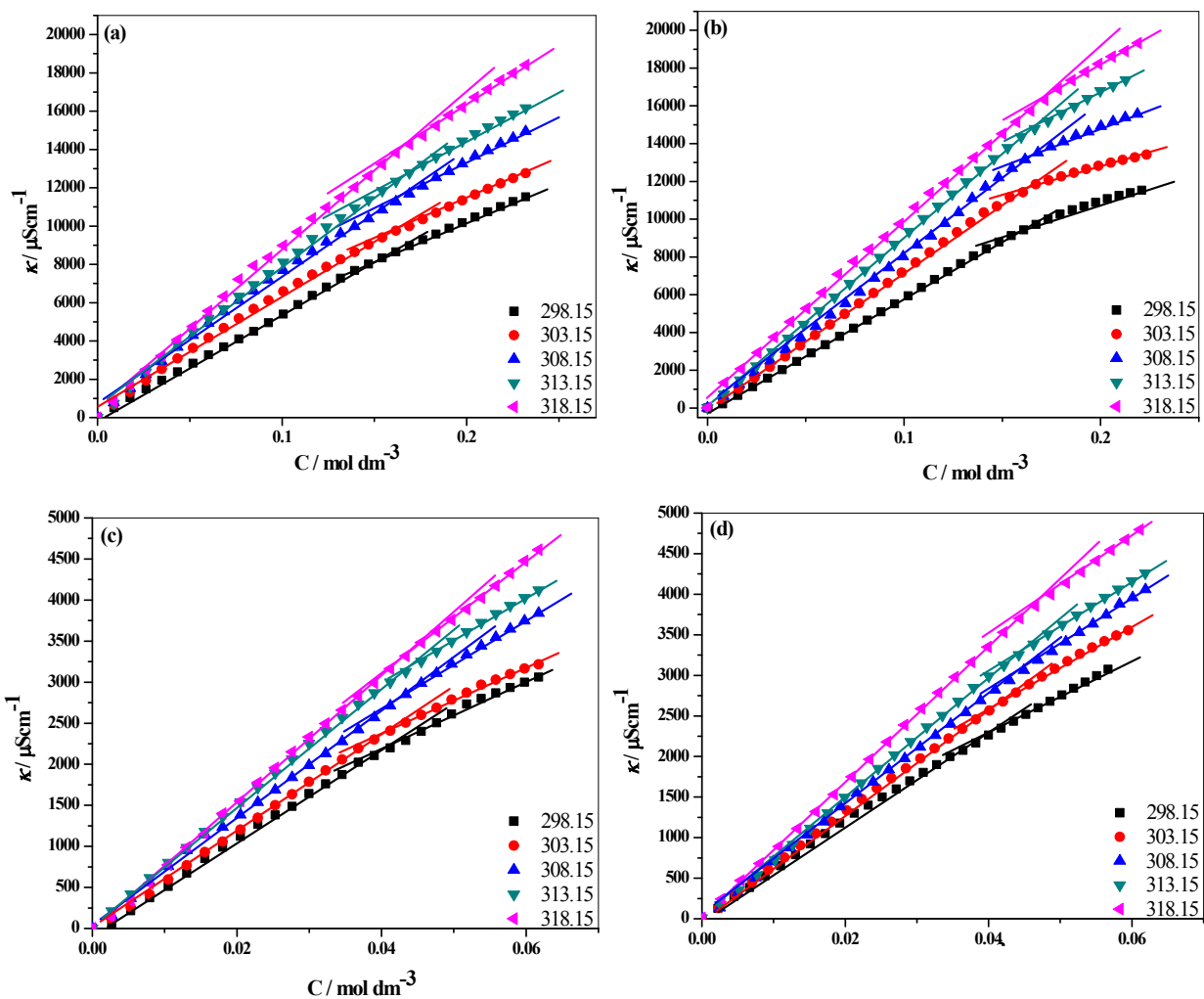


Fig. S1. Variation of specific conductivity (κ) as a function of concentration of $[\text{C}_8\text{mim}][\text{Cl}]$ and $[\text{C}_{10}\text{mim}][\text{Cl}]$ in the (a-c) absence; and (b-d) presence of polymer 10R5 at different temperatures.

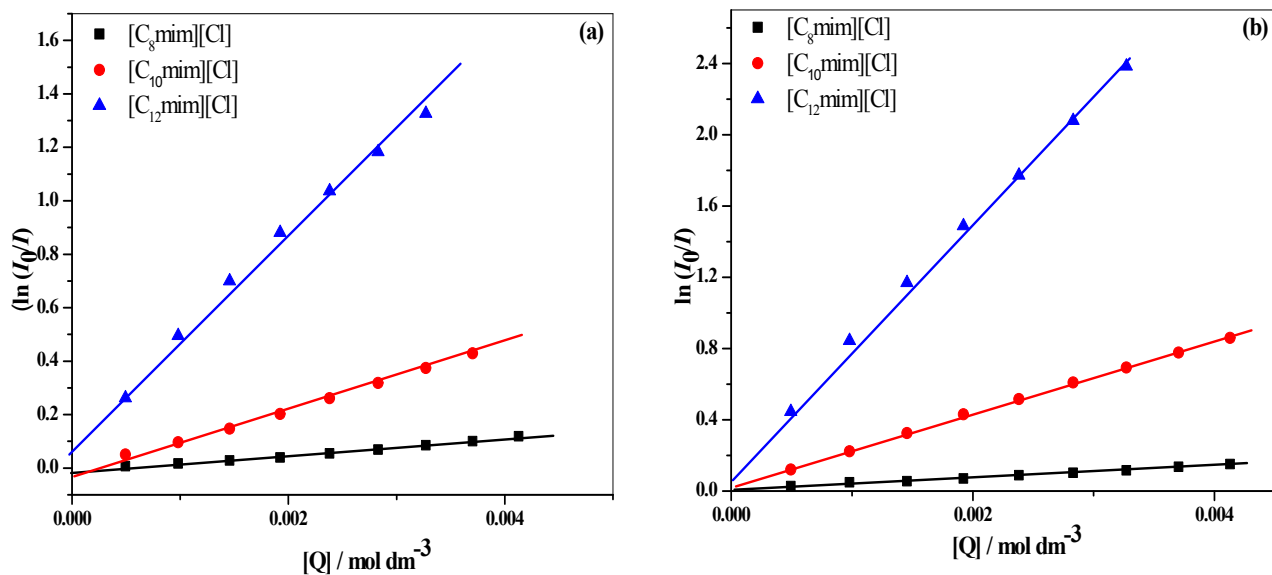


Fig. S2. Plot of $\ln(I_0/I)$ versus $[Q]$ for 10R5- $[C_8\text{mim}][\text{Cl}]$, $[C_{10}\text{mim}][\text{Cl}]$ and $[C_{12}\text{mim}][\text{Cl}]$ (a) in the absence and (b) in the presence of polymer.

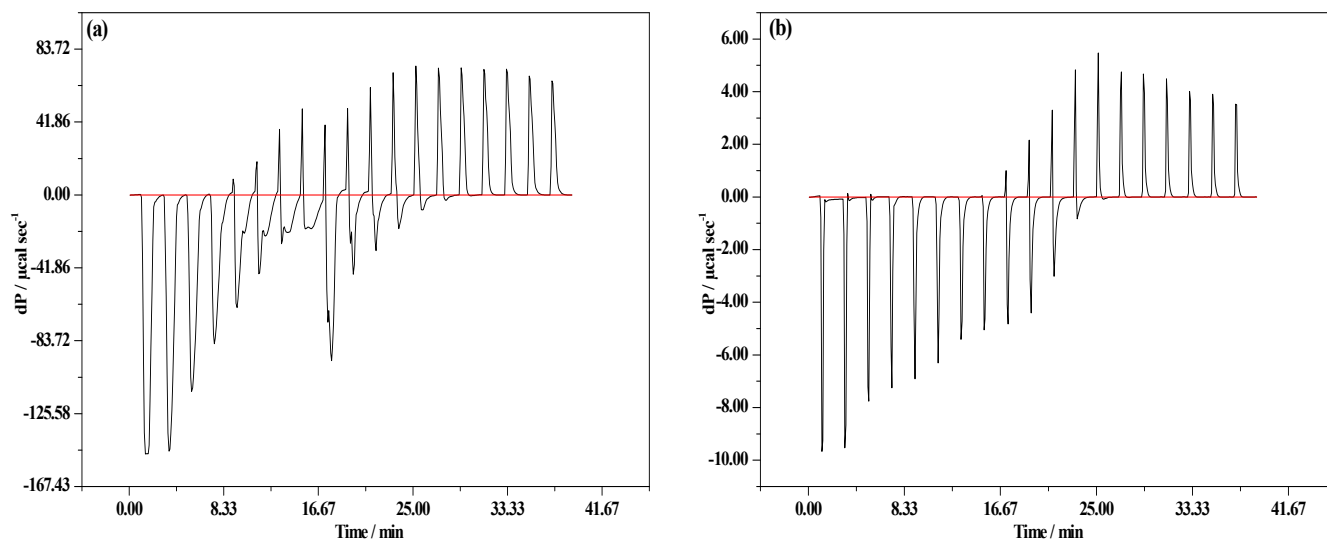


Fig. S3. Variation of differential power (dP) versus time for aqueous solution of (a) $[\text{C}_{10}\text{mim}][\text{Cl}]$ (b) $[\text{C}_{12}\text{mim}][\text{Cl}]$.

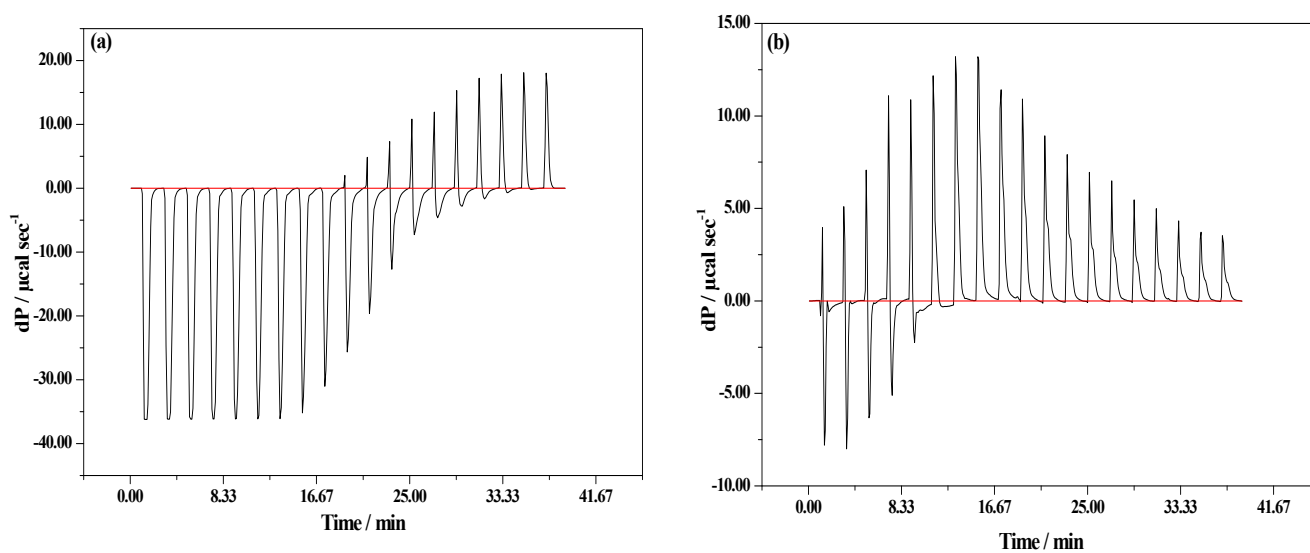


Fig. S4. Variation of differential power (dP) versus time for the interaction of 10R5 with (a) $[\text{C}_{10}\text{mim}][\text{Cl}]$ (b) $[\text{C}_{12}\text{mim}][\text{Cl}]$.

Table S1. Hydrodynamic diameter (D_h) and polydispersity (P) as a function of SAILs concentration in 10R5-[C₈mim][Cl] [C₁₀mim][Cl], [C₁₂mim][Cl] mixed systems.

conc. (mmol dm ⁻³)	D_h (nm)	P	conc. (mmol dm ⁻³)	D_h (nm)	P	conc. (mmol dm ⁻³)	D_h (nm)	P
10R5-[C ₈ mim][Cl]			10R5-[C ₁₀ mim][Cl]			10R5-[C ₁₂ mim][Cl]		
0.0	3.1	0.24	0.0	3.1	0.24	0.0	3.1	0.24
2.5	3.1	0.34	1.0	68.0	0.39	1.0	30.0	0.34
7.4	3.1	0.44	2.0	78.0	0.41	2.0	55.0	0.14
12.2	22.0	0.37	4.3	78.0	0.46	3.4	78.0	0.10
23.8	68.0	0.31	6.5	78.0	0.48	4.8	84.0	0.13
34.9	141.0	0.21	9.9	85.0	0.59	7.0	91.0	0.17
51.6	154.0	0.18	13.0	91.0	0.65	9.1	13.0	0.19
67.1	164.0	0.14	16.7	105.0	0.59	11.5	16.0	0.22
85.1	164.0	0.22	20.0	115.0	0.65	13.8	18.0	0.24
101.6	164.0	0.22	25.9	122.0	0.62	16.0	18.0	0.26
131.0	78.0	0.25	31.0	105.0	0.61	18.0	21.0	0.27
156.4	78.0	0.28	35.5	91.0	0.60	20.0	21.0	0.27
178.5	91.0	0.28	39.4	91.0	0.63			
206.7	105.0	0.30	42.9	91.0	0.60			
230.5	122.0	0.35						

References

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3. P. Mukherjee, The nature of the association equilibria and hydrophobic bonding in aqueous solutions of association colloids, Adv. Colloid Interface Sci. 1 (1967) 242-275.