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Micelle formation by *N*-alkyl-*N*-methylpiperidinium bromide ionic liquids in aqueous solution

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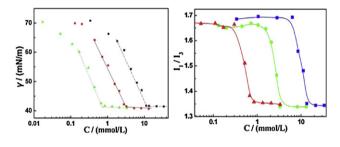
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HIGHLIGHTS

G R A P H I C A L A B S T R A C T

- Micelles are formed by N-alkyl-N-methylpiperidinium (C_nPDB) bromides in aqueous solution.
- C_nPDB surface activity is compared with other surfactants of N-heterocycle headgroups.
- Micelle formation is entropy- or enthalpy-driven at low or high temperatures.

The micelle formation by the surface active ionic liquids (ILs) *N*-alkyl-*N*-methylpiperidinium bromide $C_n PDB$ (n = 12, 14, 16) in aqueous solution is investigated through the characterizations of surface tension, electrical conductivity, and steady-state fluorescence measurements.



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ABSTRACT

The micelle formation by the surface active ionic liquids (ILs) *N*-alkyl-*N*-methylpiperidinium bromide C_n PDB (n = 12, 14, 16) in aqueous solution has been investigated through the characterizations of surface tension, electrical conductivity and steady-state fluorescence measurements. From the curves of surface tension vs C_n PDB concentration at 25 °C, the critical micelle concentration (cmc), the effectiveness of the surface tension reduction (Π_{cmc}), the maximum surface excess concentration (Γ_{max}) and the minimum area (A_{min}) occupied per surfactant molecule at the air/water interface are calculated. Through the electrical conductivity measurements, the cmc values at different temperatures and a series of thermodynamic parameters ($\Delta G_m^{\circ}, \Delta H_m^{\circ}$ and ΔS_m°) for the formation of micelles are evaluated in the temperature range of 25–45 °C. The steady-state fluorescence measurement is also employed to investigate the micelle formation of C_n PDB.

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1. Introduction

Ionic liquids (ILs), a class of fascinating organic molten salts at ambient temperature, are the focus of many scientific researchers due to their extraordinary physicochemical properties, such as high ionic conductivity, negligible vapor pressure, nonflammability, low melting temperature, and wide electrochemical window [1]. They have therefore exhibited many potential applications in the areas of catalysis, electrochemistry, preparation of novel nanomaterials, organic synthesis, liquid/liquid extraction, and green chemistry [2–5].

The great advantage of ILs is their structural designability. Their structures can be tailored by changing the cations (imidazolium, pyridinium, and pyrrolidinium) or anions (Br⁻, Cl⁻, BF₄⁻, PF₆⁻, CH₃COO⁻, CF₃COO⁻, CF₃SO₃⁻, and (CF₃SO₂)₂N⁻), which are all referred in the previous studies [6–11]. Some ILs with novel structures are synthesized and show many interesting properties. Among them, the surface active ILs (SAILs) have been developed and attracted much attention [12]. Similar to the ionic surfactant, the SAILs also have a hydrophobic chain and a hydrophilic

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