



Cite this: *Phys. Chem. Chem. Phys.*,
2015, 17, 22170

Effects of constituent ions of a phosphonium-based ionic liquid on molecular organization of H_2O as probed by 1-propanol: tetrabutylphosphonium and trifluoroacetate ions[†]

Takeshi Morita,^{*a} Kumiko Miki,^b Ayako Nitta,^a Hiroyo Ohgi^a and Peter Westh^c

Aqueous solutions of tetrabutylphosphonium trifluoroacetate, $[\text{P}_{4444}]\text{CF}_3\text{COO}$, exhibit a liquid–liquid phase transition with a lower critical solution temperature. Herein, we characterized the constituent ions, $[\text{P}_{4444}]^+$ and CF_3COO^- , in terms of their effects on the molecular organization of H_2O on the basis of 1-propanol probing methodology devised by Koga *et al.* The resulting characterization of the hydrophobicity/hydrophilicity is displayed on a two-dimensional map together with previous results, for a total of four cations and nine anions of typical ionic liquid (IL) constituents. The results indicate that $[\text{P}_{4444}]^+$ is the most significant amphiphile with strong hydrophobic and equally strong hydrophilic contributions among the group of constituent cations of ILs studied so far. The hydration number for $[\text{P}_{4444}]^+$ was evaluated to be $n_{\text{H}} = 72$, which is three times larger than that of a typical imidazolium-based cation, $[\text{C}_4\text{mim}]^+$. Self-aggregation of $[\text{P}_{4444}]^+$ was found to occur in an aqueous solution of $[\text{P}_{4444}]\text{CF}_3\text{COO}$ above 0.0080 mole fraction of the IL.

Received 21st April 2015,
Accepted 21st July 2015

DOI: 10.1039/c5cp02329g

www.rsc.org/pccp

Introduction

Ionic liquids (ILs) have gained considerable attention as they have radically changed the concept of the nature of liquids.^{1–5} ILs consisting entirely of ions have been increasingly proposed as alternatives to conventional organic solvents. Recent studies have focused on mixed systems composed of ILs and other molecular liquids, rather than neat ILs, for use as functional materials.^{6–12} In particular, thermoresponsive phase behaviour of aqueous solutions of ILs has been extensively investigated in relation to phase transitions with lower critical solution temperatures (LCSTs)^{6,12,13} as well as upper critical solution temperatures (UCSTs).^{13a,c,d,m,14}

A UCST behaviour is understandable; a strong attraction in terms of enthalpy between species would cause phase separation at low temperatures, while at higher temperatures the total entropic effect drives the system to a random mixture. This transition is caused because the mixing entropic contribution in the mixing Gibbs energy generally becomes more dominant as the temperature (T) increases. Since the enthalpy–entropy

compensation is operative particularly in aqueous solutions, the temperature could tip the balance of $\Delta_{\text{mix}}H$ and $T\Delta_{\text{mix}}S$ terms, where $\Delta_{\text{mix}}H$ and $\Delta_{\text{mix}}S$ are the change of the mixing enthalpy and entropy, respectively. However, an LCST behaviour is not as simple to explain. The present aqueous solution of $[\text{P}_{4444}]\text{CF}_3\text{COO}$ exhibits LCST-type phase behaviour with the critical point at 29.2 °C and 0.025 mole fraction of the IL.⁷ The chemical structure of $[\text{P}_{4444}]\text{CF}_3\text{COO}$ is shown in Fig. 1.

It has been suggested that LCST behaviour depends on the balance of hydrophilicity and hydrophobicity of the IL constituent ions. Kohno *et al.*¹⁵ reported that LCST-type phase behaviours of aqueous solutions of ILs strongly depend on the “hydrophilicity” of each constituent cation and anion. Their analysis was based on a one-dimensional scale with hydrophilicity and hydrophobicity at the extreme ends. The miscibility of imidazolium-based ILs with

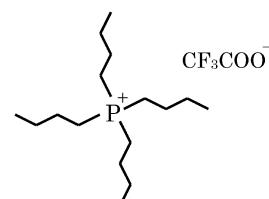


Fig. 1 Chemical structure of tetrabutylphosphonium trifluoroacetate, $[\text{P}_{4444}]\text{CF}_3\text{COO}$. Aqueous solution of the IL exhibits an LCST-type phase transition. In the present study, $[\text{P}_{4444}]\text{Cl}$ and NaCF_3COO were used for characterization of the individual constituent ions, $[\text{P}_{4444}]^+$ and CF_3COO^- , respectively.

^a Graduate School of Advanced Integration Science, Chiba University, Chiba 263-8522, Japan. E-mail: moritat@faculty.chiba-u.jp

^b Department of Liberal Arts and Basic Sciences, College of Industrial Technology, Nihon University, Narashino, Chiba 275-8575, Japan

^c NSM Research for Functional Biomaterials, Roskilde University, Roskilde DK-4000, Denmark

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c5cp02329g



water was evaluated from estimated partition coefficients in octanol–water systems.^{16,17} The origin of LCST- and UCST-type phase transitions of mixtures of thiophene with two ILs, [C₄C₁mim]SCN and [C₄C₁mim][NTf₂], was investigated using NMR spectroscopy and molecular dynamics simulations by Batista *et al.*¹⁸

Koga *et al.*^{19–21} devised a method by which an individual ion can be characterized in terms of a two-dimensional (2D) hydrophilicity and hydrophobicity scale. This technique is known as the 1-propanol (1P)-probing methodology. By this method, “amphiphiles” with hydrophobic and hydrophilic contributions can be quantitatively assessed. Here, we applied this method to characterize the constituent ions of [P₄₄₄₄]CF₃COO to seek a deeper insight.

A differential thermodynamic approach to characterize the effects of a solute on the molecular organization of H₂O was devised by Koga *et al.*¹⁹ They realized that aqueous solutions consist of three composition regions in each of which the mixing scenario on the molecular level is qualitatively different. The mixing scenario is identified by the term “mixing scheme” instead of solution structure, since the word “structure” implies a stable, ordered molecular arrangement. The mixing schemes are labelled as I, II or III from the H₂O-rich regions. In Mixing Scheme I, the hydrogen bonds of H₂O are bond-percolated throughout the entire bulk of H₂O. The transition to Mixing Scheme II from I is regarded as a loss of bond percolation when the hydrogen bond probability is reduced to the percolation threshold. The 1P-probing methodology is applicable only in the region of Mixing Scheme I.

Although the 1P-probing methodology was described in detail elsewhere,^{19,20} a brief description is given here. This methodology is based on the finding that, within Mixing Scheme I, the effects of separate solutes on H₂O are additive.¹⁹ Thus, we study a ternary system, 1P–S–H₂O, where S is the test sample whose effect on H₂O is being examined. In the ternary system of 1P–S–H₂O, the thermodynamic signature of 1P, H_{1P1P}^E (defined below), is evaluated. Modification to its x_{1P} -dependence pattern due to the presence of S is used to characterize the effect of S on H₂O. H_{1P1P}^E is the enthalpic 1P–1P interaction in a complex ternary system and is defined as,

$$H_{1P1P}^E \equiv N(\partial H_{1P}^E / \partial n_{1P}) = (1 - x_{1P})(\partial H_{1P}^E / \partial x_{1P}), \quad (1)$$

where

$$H_{1P}^E \equiv (\partial H^E / \partial n_{1P}), \quad (2)$$

and H^E is the total excess enthalpy in the (p , T , n_{1P} , n_S , n_W) variable system, $N = n_{1P} + n_S + n_W$ and $x_{1P} = n_{1P}/N$. The subscripts 1P, S and W signify 1-propanol, test sample S and H₂O, respectively. We directly measure the excess partial molar enthalpy of 1P, H_{1P}^E . To do so, we perturb the ternary system by increasing n_{1P} to $n_{1P} + \delta n_{1P}$, with a fixed initial mole fraction of S, $x_S^0 = n_S/(n_S + n_W)$, and measure the H^E response, δH^E . We approximate the quotient $\delta H^E / \delta n_{1P}$ to the partial derivative in eqn (2), and evaluate H_{1P1P}^E by differentiating the data set (x_{1P} , H_{1P}^E) using the far right of eqn (1).

Fig. 2 shows a schematic representation of the changes in the H_{1P1P}^E pattern induced by the presence of various classes of solute, S. As shown in Fig. 2(a)–(d), the peak top is named point X, which marks the end of Mixing Scheme I. Upon addition of S, point X shifts depending on the nature of S. The shifts are in general linear to the initial mole fraction of S, x_S^0 , within Mixing Scheme I. The slope of the westward shift (*i.e.* to the negative direction of x_{1P}) of point X per unit increase in x_S^0 is taken as the *hydrophobicity index*, while that towards the south (*i.e.* to the negative direction of the H_{1P1P}^E axis) is the *hydrophilicity index*.

Fig. 2(a) shows a typical change in H_{1P1P}^E for hydration centres such as in Na⁺ and Cl[–] ion pairs.^{22,23} Upon addition of the salt, the H_{1P1P}^E pattern is compressed to the west. This compression indicates that, upon addition of the Na⁺ and Cl[–] ion pair, the available H₂O molecules for 1P to interact with are reduced. We thus interpret this westward shift to be due to hydration of the ions. From the westward shift of point X as a function of x_S^0 , the hydration number, n_H , can be evaluated for Na⁺ and Cl[–]. It must be stressed that at the starting point, $x_{1P} = 0$, the value of H_{1P1P}^E remains the same even in the presence of Na⁺ and Cl[–]. This invariance and the fact that the value of H_{1P1P}^E at point X also remains the same led to the following suggestions: (1) Na⁺ and Cl[–] ions are hydrated by a number of H₂O molecules, (2) the hydrating H₂O molecules are unavailable to interact with 1P and (3) the bulk H₂O molecules away from the hydration shells are unperturbed by Na⁺ and Cl[–] even in the presence of the ions. This finding can be utilized to characterize individual ions. For a given test ion, we chose the counter ion Na⁺ or Cl[–] and applied the 1P probing on the combined salt. Namely, we used tetrabutylphosphonium chloride, [P₄₄₄₄]Cl, and sodium trifluoroacetate, NaCF₃COO, to characterize [P₄₄₄₄]⁺ and CF₃COO[–], respectively.

Fig. 2(b) shows the behaviour of the hydrophobic solute. For a solute almost equally hydrophobic as 1P such as 2-propanol²⁴ the H_{1P1P}^E pattern shifts parallel to the west, as shown in the figure. This parallel shift is expected when compared to the case in which 1P was added as S to x_S^0 (*i.e.* S = 1P). For a hydrophobe that is stronger (or weaker) than the probing 1P, such as *tert*-butanol²⁵ (or ethanol²⁶), the westward shift is greater (or lesser), reflecting a larger (or smaller) n_H . This westward shift of point X indicates that a hydrophobe is hydrated by H₂O molecules, making them unavailable to interact with 1P, as was the case for the hydration centre. In addition, the value of H_{1P1P}^E at point X shifts northward (or southward), reflecting the fact that the ability of *tert*-butanol (or ethanol) to reduce the hydrogen bond probability of bulk H₂O away from the hydration shell is greater (or lesser) than 1P, as studied earlier.¹⁹ Furthermore, the hydrogen bonding within the hydration shells is more organized than it is in bulk H₂O.

Fig. 2(c) shows the pattern change for the hydrophilic solute. This finding, as well as others,¹⁹ led to the interpretation that hydrophiles form hydrogen bonds directly to the existing hydrogen bond network of H₂O and maintain the hydrogen bond connectivity of the network; thus, they act as impurities in the network. As such, they break the H donor/acceptor symmetry. Hence, the southward shift apparent in the figure is



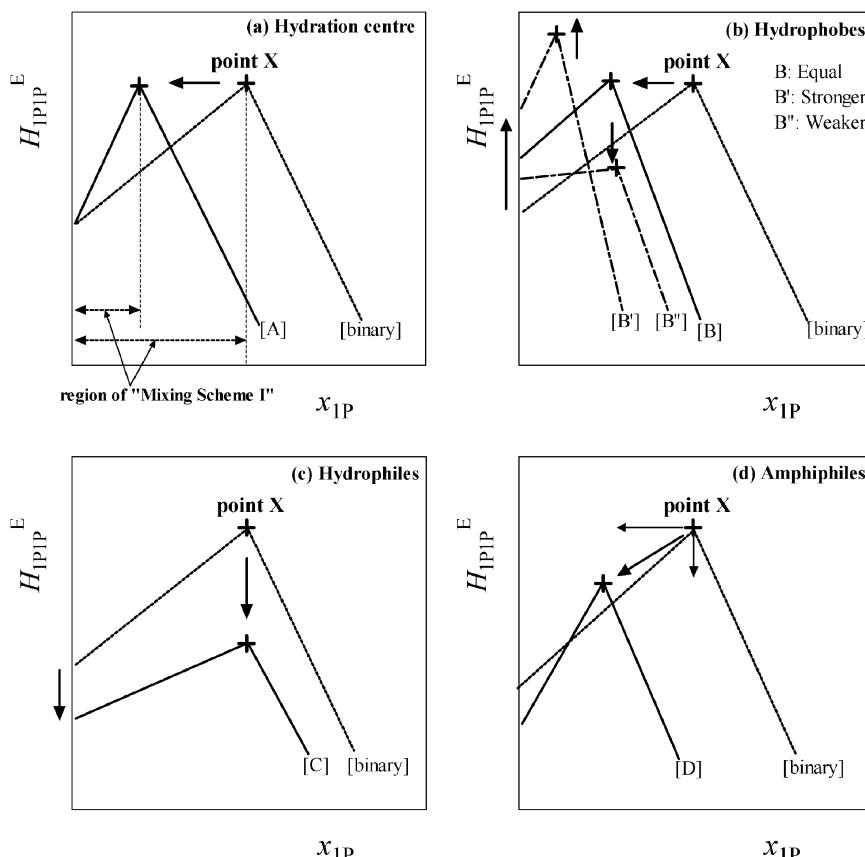


Fig. 2 Induced changes of H_{1P1P}^E pattern in the presence of various types of solute S. [binary] represents the binary 1P–H₂O system without test solute S. (a) Effects of the hydration centre on the pattern per unit increase in test sample S. Region of Mixing Scheme I is defined as $x_{1P} = 0$ to x_{1P} at point X, as described on the horizontal axis, (b) the effects of hydrophobes, (c) hydrophiles and (d) amphiphiles.

interpreted as a reduction of the net entropy-volume cross fluctuation.

Fig. 2(d) is for the amphiphilic solute. The effects seem to be a combination of those observed for hydrophobic and hydrophilic moieties. Their westward and southward components show contributions from hydrophobic and hydrophilic moieties, respectively. Typical IL constituent ions generally show amphiphilic responses to 1P-probing studies with strong hydrophobic and equally strong hydrophilic characteristics.^{27–30} These results fit into the special properties of ILs. Low melting points for ionic compounds can be related to the strength of hydrophobicity and/or hydrophilicity.^{20,30}

With this pair of coordinates, the characterization of each species, including individual constituent ions, is displayed on a 2D map of hydrophobicity/hydrophilicity.

Experimental

Titration calorimetry for 1P probing

Tetrabutylphosphonium chloride, [P₄₄₄₄]Cl, (Aldrich, >98%) was used as the sample for characterization of the cation, [P₄₄₄₄]⁺. Sodium trifluoroacetate, NaCF₃COO, (Sigma-Aldrich, >96%) was used as the sample for that of CF₃COO[−]. Deionized

H₂O was prepared using a Milli-Q system (Millipore), with a resulting resistivity of 18.2 MΩ cm. Stock solutions were prepared using the purified H₂O and the salts from freshly opened bottles. 1-Propanol (1P) (Sigma-Aldrich, Chromasolv for HPLC 99.9%) was used as supplied. Due care was exercised not to contaminate the 1P with moisture from the atmosphere. The excess partial molar enthalpy, H_{1P}^E , was determined using a TAM III semi-isothermal titration calorimeter (TA Instruments, New Castle, USA) at 25.0 °C ± 0.0001 °C and in the dynamic correction mode.^{31–33} A 1 mL stainless steel container was used as the cell. The initial volume of the sample solution was set at 750 μL and the exact amount was determined gravimetrically. The volume and duration of each titration of 1P were set at 10 μL and 10 s, respectively. The injection volume per titration was calibrated to 9.9483 μL and was used to calculate the concentration. The ratio of titrant over titrand was on the order of 10^{−2} and was well within the acceptable range for approximation of the differentiation. The interval of injection was set at 30 min. The averaged uncertainty of the titration measurements was estimated to be ±0.03 kJ mol^{−1}.

Electric conductivity

The IL, [P₄₄₄₄]CF₃COO, was synthesized and used for conductivity measurements. It was prepared by direct neutralization of



aqueous tetrabutylphosphonium hydroxide solutions with trifluoroacetic acid. After evaporation, the product was added to a dichloromethane/water biphasic system and the resulting mixture was washed several times with distilled water. The required $[P_{4444}]CF_3COO$ was dissolved in dichloromethane phase, and was dried in vacuum for 24 h at 60 °C prior to use. The structure and purity were confirmed by 1H NMR spectroscopy. The procedure of the synthesis has been described elsewhere in detail.⁷

The electric conductivity of the aqueous solution was measured using an electrical conduction metre with automatic temperature compensation (Hanna instruments, DiST4). The measured concentration range was from 0.001127 to 0.04610 mole fraction of the IL.

Results and discussion

The data of H_{1P}^E are deposited as Table S1 of the ESI,† and are plotted in Fig. 3. A sigmoidal increase in H_{1P}^E with an inflection point is evident as long as x_S^0 is appropriate for performing the 1P-probing methodology. As shown in Fig. 3, the x_{1P} -dependence pattern of H_{1P}^E changes more rapidly for $S = [P_{4444}]Cl$ with increasing x_S^0 than it does for $S = NaCF_3COO$. The inflection point seems to disappear at $x_S^0 = 0.011$ for $S = [P_{4444}]Cl$ and at $x_S^0 = 0.055$ for $S = NaCF_3COO$. To see this

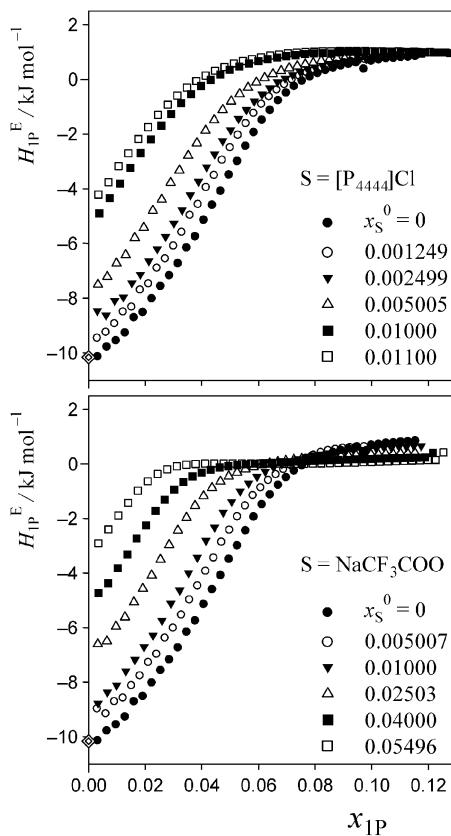


Fig. 3 Excess partial molar enthalpy, H_{1P}^E , of 1P in 1P–S–H₂O at 25 °C and given initial salt concentration, x_S^0 . Diamonds represent H_{1P}^E in dilute solution, as reported by Wadso *et al.*³⁵

behaviour more clearly, we took the derivative of H_{1P}^E with respect to x_{1P} .

Evaluation of H_{1P1P}^E from H_{1P}^E data

With a good set of raw H_{1P}^E data with an averaged uncertainty of ± 0.03 kJ mol⁻¹ and small increments in x_{1P} , the raw H_{1P}^E data themselves were used to calculate the derivative on the far right of eqn (1). In general, a numerical derivative directly using two neighbouring data points causes large noise, especially in applications with raw experimental data. In the present study, we originally utilized a differential method based on eqn (3) for calculation of the partial molar quantities with higher-order derivatives. We approximate the slope of the tangent at the i -th point as the weighted average of the slopes of the two adjacent arcs; one between the $(i - 1)$ -th and i -th data points and another between the i -th and $(i + 1)$ -th data points.

$$H_{1P1P}^E(i) = \{1 - x_{1P}(i)\} \left\{ \frac{\partial H_{1P}^E(i)}{\partial x_{1P}(i)} \right\} \approx \frac{\{1 - x_{1P}(i)\}}{\{x_{1P}(i+1) - x_{1P}(i-1)\}} \left[\frac{\{H_{1P}^E(i+1) - H_{1P}^E(i)\}}{\{x_{1P}(i+1) - x_{1P}(i)\}} \{x_{1P}(i) - x_{1P}(i-1)\} \right. \\ \left. + \frac{\{H_{1P}^E(i) - H_{1P}^E(i-1)\}}{\{x_{1P}(i) - x_{1P}(i-1)\}} \{x_{1P}(i+1) - x_{1P}(i)\} \right] \quad (3)$$

The x_{1P} interval of $x_{1P}(i)$ to $x_{1P}(i + 1)$, δx_{1P} , was approximately 0.003 in the present measurements. It is found that $\delta x_{1P} \approx 0.008$ in graphical evaluation normally exercised is appropriate to approximate the derivative of eqn (1) with the quotient $\delta H_{1P}^E/\delta n_{1P}$.³⁴ In the present calculations, the total δx_{1P} for three neighbouring data points was approximately 0.006.

Certainty in evaluation of the partial molar quantities with higher-order derivatives is of particular importance in the present study. Fig. 4(a)–(c) show comparisons of H_{1P1P}^E determined using eqn (3) and that determined using other evaluation methods. Fig. 4(a) shows comparison between the H_{1P1P}^E pattern obtained using eqn (3) and that evaluated by the graphical method normally exercised. As shown in Fig. 4(b), combination of eqn (3) and the graphical evaluation shows improvement especially in the pre-peak region. The point X evaluated by the H_{1P1P}^E pattern corresponds to each other regardless of the method. Fig. 4(c) shows comparison between eqn (3) and simple derivative using two neighbouring data points. As mentioned above, the calculation using eqn (3) is superior to the simple derivative using two alternating data points with regard to noise reduction, as shown in Fig. 4(c). Therefore, it is concluded that the calculation based on eqn (3) is the most certain for evaluation of partial molar quantities with higher-order derivatives in the present data set. Thus, calculation based on eqn (3) was utilized here for evaluation of H_{1P1P}^E , leading to determination of the hydrophobicity and hydrophilicity indices.

Hydrophobicity/hydrophilicity obtained from the H_{1P1P}^E pattern

Fig. 5 and 6 show the patterns of resulting H_{1P1P}^E for $S = [P_{4444}]Cl$ and $S = NaCF_3COO$, respectively. The peak top of H_{1P1P}^E appears

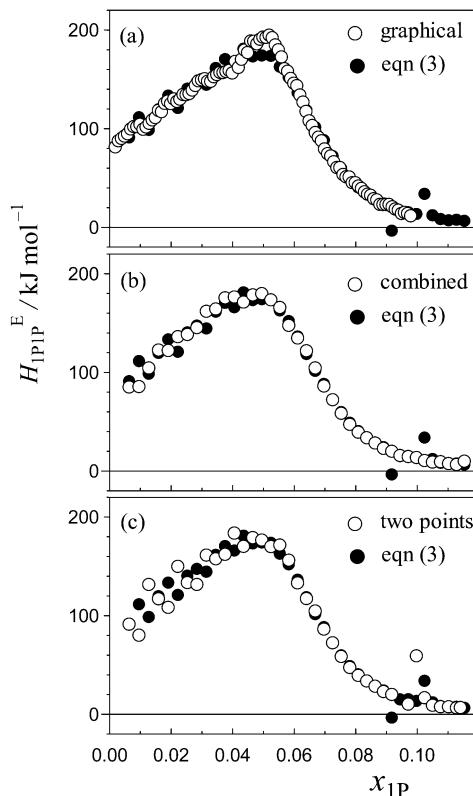


Fig. 4 Comparison of H_{1P1P}^E pattern for the binary (1-propanol–H₂O) mixture calculated based on differentiation using eqn (3) (solid circles) with those using other evaluation methods (open circles); (a) typical graphical evaluation normally exercised for 1P-probing studies,³⁴ (b) combination of calculation using eqn (3) and graphical evaluation, (c) simple derivative using two data points with $\delta x_{1P} \approx 0.006$.

at 0.049 of x_{1P} for binary (1-propanol–H₂O) mixture.¹⁹ The peak top shifts to the west (direction of smaller x_{1P}) as x_S^0 increases. Point X nearly disappears at $x_S^0 = 0.0110$ for S = [P₄₄₄₄]Cl and $x_S^0 = 0.0550$ for S = NaCF₃COO, qualitatively indicating that the hydrophobicity of [P₄₄₄₄]Cl is much larger than that of NaCF₃COO. The starting point of H_{1P1P}^E at $x_{1P} = 0$ shifts to the north (direction of larger H_{1P1P}^E) with increasing x_S^0 . This northward shift also corresponds to the strong hydrophobicity of [P₄₄₄₄]⁺. For NaCF₃COO, the equivalent shift at $x_{1P} = 0$ is lesser.

The degrees of the shifts are plotted in Fig. 7 and 8. The slopes of x_{1P} loci shown in Fig. 7(a) and 8(a) yielded the hydrophobicity indices and hydration numbers, n_H . The contribution from counter ions (Na⁺ or Cl⁻) was corrected by subtraction in the evaluation. The slopes of the H_{1P1P}^E loci in Fig. 7(b) and 8(b) directly correspond to the hydrophilicity indices in the present evaluation. The hydrophilicity indices of the counter ions of Na⁺ and Cl⁻ used in the present study are equal to zero, since these ions are classified as the hydration centre. As mentioned in Introduction, hydration centres such as Na⁺ and Cl⁻ unperturb the bulk H₂O molecules away from the hydration shells around the ions. Hydrophobicity/hydrophilicity indices and hydration numbers of [P₄₄₄₄]⁺ and CF₃COO⁻ are listed in Table 1.

The results indicate that [P₄₄₄₄]⁺ is an amphiphile with strong hydrophobic and equally strong hydrophilic contributions.

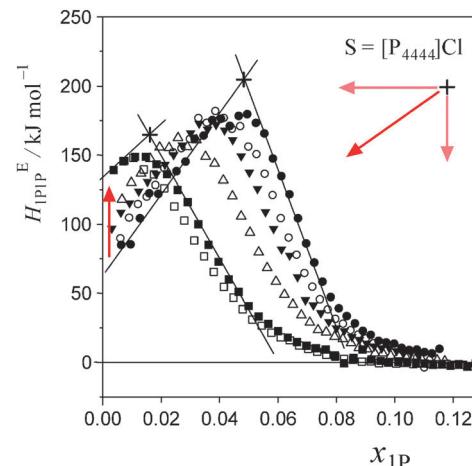


Fig. 5 Enthalpic 1P–1P interaction, H_{1P1P}^E , of 1P–[P₄₄₄₄]Cl–H₂O at 25 °C for given initial salt concentrations, x_S^0 . The initial concentrations, x_S^0 , were set at 0, 0.001249, 0.002499, 0.005005, 0.01000 and 0.01100 from right to left in the figure. Mixing Scheme I is classified from $x_{1P} = 0$ to x_{1P} at point X. The region decreases with increasing x_S^0 and nearly disappears at $x_S^0 \approx 0.011$. Point X is largely compressed in the westward and southward directions.

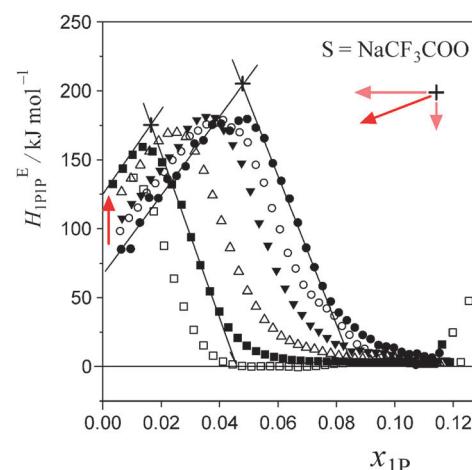


Fig. 6 Enthalpic 1P–1P interaction, H_{1P1P}^E , of 1P–NaCF₃COO–H₂O at 25 °C for given initial salt concentrations, x_S^0 . The initial concentrations, x_S^0 , were set at 0, 0.005007, 0.01000, 0.02503, 0.04000 and 0.05496 from right to left in the figure. Mixing Scheme I decreases with increasing x_S^0 and disappears around $x_S^0 \approx 0.055$. Point X is compressed mainly in the westward direction.

[P₄₄₄₄]⁺ forms a large hydration shell with a hydration number of $n_H = 72$, which is three times larger than that of typical imidazolium-based cation 1-butyl-3-methylimidazolium, [C₄mim]⁺.²⁷ The value of a hydration number for CF₃COO⁻ was found to be $n_H = 10$. On the basis of earlier findings in 1P-probing studies on a series of carboxylates,³⁶ one H₂O molecule out of the 10 H₂O molecules should be used for hydration of the –COO⁻ side and the remaining 9.0 H₂O molecules for the fluoroalkyl group, –CF₃. This hydration number is much larger than that of the alkyl group in CH₃COO⁻; the hydration number for –CH₃ in CH₃COO⁻ is estimated to be 2.7.³⁶ The fact that the fluoroalkyl

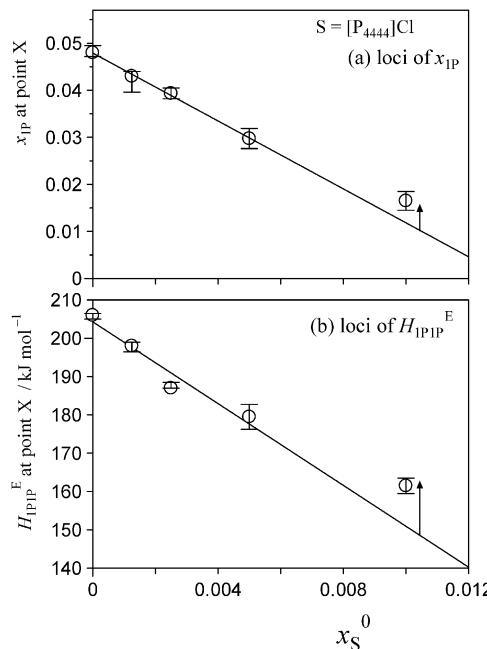


Fig. 7 Loci of point X as a function of the initial salt concentration, x_S^0 , for $S = [P_{4444}]Cl$. (a) The loci of x_{1P} at 25 °C. The slope gives the hydrophobicity index and hydration number, n_H . These parameters were evaluated from the slope following subtraction of the Cl^- counterion contribution. (b) The loci of H_{1P1P}^E at 25 °C. The slope corresponds to the hydrophilicity index, as Cl^- is classified as the hydration centre. The hydrophilicity index of Cl^- is zero.

group is a stronger hydrophobe than the alkyl group could be related to this finding;³⁷ an aqueous solution of the CF_3COO^- salt of $[P_{4444}]^+$ shows LCST, and that of the CH_3COO^- salt does not.

Self-aggregation behaviour of $[P_{4444}]^+$

As shown in Fig. 7, the induced changes at $x_S^0 = 0.010$ for $S = [P_{4444}]Cl$ depart from the linear relationship defined in the lower x_S^0 region. This departure shifts to the upper side beyond the data error in the present evaluation. This departure hints the self-aggregation of the kind found in imidazolium-based ILs. For $[C_4C_1mim]Cl$ ²⁹ and $[C_4mim]Cl$,²⁷ similar plots are seen that show a break in the slopes at $x_S^0 = 0.006$ and 0.014, respectively. For aqueous solutions of $[C_4mim]BF_4$, this similar behaviour was interpreted as self-aggregation of cations through NMR spectroscopy,³⁸ electric conductivity,^{39–41} small-angle neutron scattering,^{41,42} density,⁴³ speed of sound⁴³ and surface tension.^{39,41} Thus, we suggest that departure of the induced change at $x_S^0 = 0.010$ for the case shown in Fig. 7 could also be caused by self-aggregation of $[P_{4444}]^+$.

For further investigation of this issue, electric conductivity measurements were conducted for the aqueous solutions of $[P_{4444}]CF_3COO$. The obtained conductivity was converted to molar conductivity according to the Kohlrausch empirical relation.⁴⁴ Fig. 9 shows the isotherm of the molar conductivity as a function of the square root of IL concentration. The linear portion represents the usual behaviour of ionic solutions in the dilute region. This linearity seems to start to deviate at

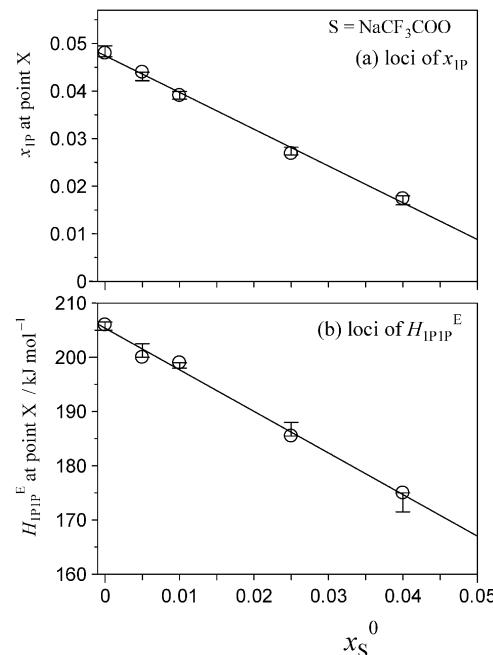


Fig. 8 Loci of point X as a function of initial salt concentration, x_S^0 , for $S = NaCF_3COO$. (a) The loci of x_{1P} at 25 °C. The slopes give the hydrophobicity index and hydration number, n_H . These parameters were evaluated from the slope following subtraction of the Na^+ counterion contribution. (b) The loci of H_{1P1P}^E at 25 °C. The slope corresponds to the hydrophilicity index, as Na^+ is classified as the hydration centre. The hydrophilicity index of Na^+ is zero.

$\sqrt{c} = 19.6$ (mmol L⁻¹)^{1/2}, which corresponds to a mole fraction of 0.0080. The fact that the conductivity shows deviation from the usual behaviour at $x_{IL} = 0.0080$ suggests that $[P_{4444}]CF_3COO$ dissociates into its constituent ions in the dilute region and aggregation of $[P_{4444}]^+$ begins at a mole fraction of approximately 0.0080, as the case for imidazolium-based ILs. Considering the larger hydrophobicity (hydration number) of $[P_{4444}]^+$ compared to those of $[C_4mim]^+$ and $[C_4C_1mim]^+$, the breakpoint (or aggregation) of $[P_{4444}]^+$ might be expected at a much lower concentration; however, this was not observed. This discrepancy could be related to the difference in the aggregation mechanisms of imidazolium-based and phosphonium-based cations. The imidazolium ions have an ionic head and alkyl tail, while phosphonium-based ions have an ionic centre and four alkyl tails extending in random directions. Wang *et al.* suggested that the aggregation aspect of $[P_{4444}]CF_3COO$ in aqueous solution can be described in terms of microemulsion-like aggregation at the mesoscale level in higher concentration ranges near the critical point.⁴⁵

2D map of hydrophobicity/hydrophilicity

Fig. 10 shows the 2D map of hydrophobicity/hydrophilicity for $[P_{4444}]^+$ and CF_3COO^- . In the map, H_2O defines the origin (0,0). 1-Propanol used as the probe is necessarily plotted at (-1,0). Relative to these points, ions with greater hydrophobic contributions spread to the west (negative horizontal direction)

Table 1 Hydrophobicity/hydrophilicity indices of $[P_{4444}]^+$ and CF_3COO^- as probed by 1-propanol

Ion	Class	Hydrophobicity	n_H^a	Hydrophilicity (kJ mol ⁻¹)	Applicable x_S^0 range	Aggregation ^b	Ref.
Cations							
$[P_{4444}]^+$	Amphiphile	-3.49	72	-5337	<0.012	$x_S^0 > 0.0080$	This work
$[C_2mim]^+$	Amphiphile	-0.39	7	-1970	<0.024	None	20, 21 and 29
$[C_4mim]^+$	Amphiphile	-1.31	26	-3227	<0.029	$x_S^0 > 0.013$	20, 21 and 27
$[C_4C_1mim]^+$	Amphiphile	-1.85	37	-6760	<0.018	$x_S^0 > 0.0060$	20, 21 and 29
Anions							
CF_3COO^-	Hydrophobe	-0.49	10	-767	<0.055	None	This work
CH_3COO^-	Hydrophobe	-0.22	3.7	0	<0.05	None	20, 21 and 36

^a Hydration number evaluated from the loci of point X. ^b Aggregation observed in the region of Mixing Scheme I.

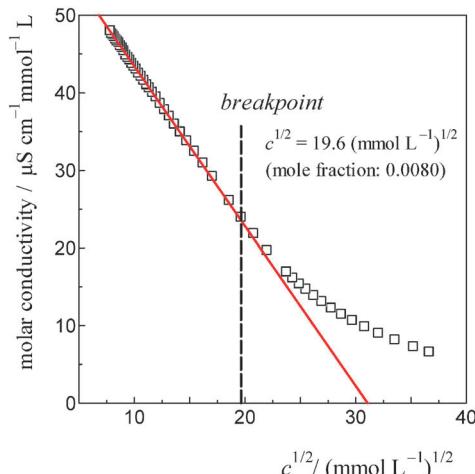


Fig. 9 The isotherm of electric conductivity for aqueous solutions of $[P_{4444}]CF_3COO$ based on the Kohlrausch empirical relation.⁴² The break point was estimated to be $c^{1/2} = 19.6$ (mmol L⁻¹)^{1/2} for $[P_{4444}]CF_3COO$, corresponding to a 0.0080 mole fraction of the IL.

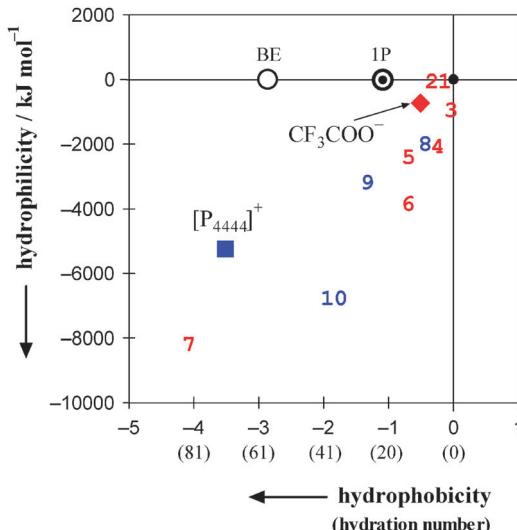


Fig. 10 2D map of hydrophobicity/hydrophilicity for typical constituent ions of ionic liquids. Values in parentheses on the horizontal axis denote hydration numbers. H_2O is located at the origin (0,0) and the probing 1P is located at (-1,0). The square and the diamond represent the present results for $[P_{4444}]^+$ and CF_3COO^- , respectively. Seven other constituent anions^{20,21,30} and three cations^{20,21} are shown together; 1: Cl^- , 2: CH_3COO^- , 3: Br^- , 4: BF_4^- , 5: $[OTf]^-$, 6: $[PF_6]^-$, 7: $[NTf_2]^-$, 8: $[C_2mim]^+$, 9: $[C_4mim]^+$, 10: $[C_4C_1mim]^+$. An aqueous solution of 2-butoxyethanol (abbreviated as BE) exhibits an LCST below 50 °C.^{50,51} See text for discussion.

and those with greater hydrophilic contributions spread to the south (negative vertical direction). A total of four cations and nine anions of typical IL constituents are plotted in Fig. 10. The cations^{20,21} are $[C_2mim]^+$, $[C_4mim]^+$, $[C_4C_1mim]^+$ and the present $[P_{4444}]^+$. The anions^{20,21,30} are Cl^- , CH_3COO^- , Br^- , BF_4^- , $[OTf]^-$, PF_6^- , $[NTf_2]^-$ and the present CF_3COO^- . As evident from Fig. 10, $[P_{4444}]^+$ shows the most significant amphiphilicity, with strong hydrophobic and equally strong hydrophilic contributions, among the constituent cations of the ILs studied. Furthermore, the IL constituent ions are located farther from the origin than are the typical inorganic small ions. Our recent study on normal ions (*i.e.* non-IL forming ions) shows that anions are located farther from the origin than cations.⁴⁶

Kohno *et al.*¹⁵ took advantage of the phase separation of IL-H₂O systems and estimated the H₂O content in the IL phase. They then determined the hydration number per mole of IL and defined it as the hydrophilicity of the IL. They concluded that the LCST behaviour strongly depended on the evaluated “hydrophilicity” indices. We point out that their hydrophilicity concept is based on the H₂O content in the IL-rich situation. In the 1P probing, hydrophilicity is only a part of the effects of the ion in question to H₂O in the water-rich region, the other part

being hydrophobicity. Saita *et al.*⁴⁷ suggested another one-dimensional scale based on salt effects of test ions fixed as $[P_{4444}]^+$ or CF_3COO^- counter ion on the LCST of $[P_{4444}]CF_3COO-H_2O$ system. It is concluded that changes in the critical temperatures of IL-H₂O systems are clearly related to the “hydrophilicity” of the target ions.

In the present 1P-probing methodology, the hydration number of the species in question is evaluated from “hydrophobicity” index. The hydrophobicity is defined on the basis of hydration of the solute with H₂O molecules, namely the formation of the hydration shell in water-rich region. Hydrophile in the 1P probing is defined as a solute that forms hydrogen bonds directly to the existing hydrogen bond network of H₂O, discussed in detail in Introduction. As shown in Fig. 10, the typical constituent ions are classified as “amphiphiles” with strong hydrophobic and equally strong hydrophilic contributions. Thus, the present



two-dimensional scale covers the ranking of “hydrophilicity” from the one-dimensional scale and might lead to a deeper understanding on the effects of molecular organization of H_2O .

Conclusions

The constituent ions of $[\text{P}_{4444}]\text{CF}_3\text{COO}$ were characterized in terms of their effects on the molecular organization of H_2O using the 1P-probing methodology. A differential method based on the weighted average of two slopes was utilized to calculate $H_{1\text{P}1\text{P}}^E$. The most significant amphiphile among the IL constituent cations studied was $[\text{P}_{4444}]^+$, which had strong hydrophobic and equally strong hydrophilic contributions. The hydration number was evaluated to be $n_{\text{H}} = 72$, which was the largest value among the group of cations. Characterization of CF_3COO^- confirmed that it was a hydrophobe with a hydration number of $n_{\text{H}} = 10$, out of which one H_2O molecule hydrated the $-\text{COO}^-$ side and the remaining 9.0 H_2O molecules the fluoroalkyl group, $-\text{CF}_3$. The self-aggregation behaviour of $[\text{P}_{4444}]^+$ in the aqueous solution of $[\text{P}_{4444}]\text{CF}_3\text{COO}$ was revealed above 0.0080 mole fraction of the IL.

It is suggested that the formation of the large hydration shell around $[\text{P}_{4444}]^+$ evaluated in the present study causes loss in excess entropy of mixing. Hence, this entropic loss must have bearing to the present LCST behaviour as mentioned in Introduction. Koga *et al.* sorted out the occurrence of phase separation with an LCST as well as a UCST in terms of signs of H_{ii}^E and S_{ii}^E , the third derivative of G^E in the binary solute(i)- H_2O system.^{19,48} Although their argument was based on a necessary but not sufficient condition, it was applied to the $[\text{C}_4\text{mim}]\text{BF}_4-\text{H}_2\text{O}$ system and the H_{ii}^E and S_{ii}^E for $i = [\text{C}_4\text{mim}]\text{BF}_4$ was found to be negative, which is appropriate for the UCST at 4 °C and $x_i = 0.07$.⁴⁹ For the present case, however, since we have not yet determined H_i^E and S_i^E for $i = [\text{P}_{4444}]\text{CF}_3\text{COO}$, we will postpone the detailed discussions about the occurrence of LCST for aqueous solution of the present IL. As shown in Fig. 10, 2-butoxyethanol (BE) was found to form an analogously large hydration shell around the molecule, $n_{\text{H}} = 58$,²³ and the aqueous solution of BE exhibits the LCST below 50 °C.^{50,51}

Acknowledgements

We would like to express deepest appreciation to Dr Y. Koga at the University of British Columbia for his discussion about the 1P-probing experiment and critical reading of the manuscript. We are deeply grateful to Prof. H. Ohno, Dr Y. Kohno and Dr S. Saita at Tokyo University of Agriculture and Technology for preparation of the ionic liquid used in this study, measurements of the electric conductivity and fruitful discussion. T.M. is grateful to Prof. K. Nishikawa for cooperation of the experiment. This work was partially supported by Grant-in-Aid for Scientific Research (JSPS KAKENHI) (No. 24550009). T.M. thanks MEXT, Japan, for Grants for Excellent Graduate Schools for the financial support for his stay at Roskilde University.

Notes and references

- 1 J. S. Wilkes and M. J. Zaworotko, *J. Chem. Soc., Chem. Commun.*, 1992, 965.
- 2 T. Welton, *Chem. Rev.*, 1999, **99**, 2071.
- 3 J. F. Wishart and E. W. Castner, Jr., “*The Physical Chemistry of Ionic Liquids*” (Editorial for Special Issue), *J. Phys. Chem. B*, 2007, **111**, 4639.
- 4 N. V. Plechkova and K. R. Seddon, *Chem. Soc. Rev.*, 2008, **37**, 123.
- 5 P. Wasserscheid and T. Welton, *Ionic Liquids in Synthesis*, Wiley-VCH, Weinheim, 2nd edn, 2008.
- 6 K. Fukumoto and H. Ohno, *Angew. Chem., Int. Ed.*, 2007, **46**, 1852.
- 7 Y. Kohno, H. Arai, S. Saita and H. Ohno, *Aust. J. Chem.*, 2011, **64**, 1560.
- 8 Y. Kohno and H. Ohno, *Chem. Commun.*, 2012, **48**, 7119.
- 9 T. V. Hoogerstraete, B. Onghena and K. Binnemans, *J. Phys. Chem. Lett.*, 2013, **4**, 1659.
- 10 A. J. L. Costa, M. R. C. Soromenho, K. Shimizu, J. M. S. S. Esperança, J. N. C. Lopes and L. P. N. Rebelo, *RSC Adv.*, 2013, **3**, 10262.
- 11 W. Li and P. Wu, *Polym. Chem.*, 2014, **5**, 5578.
- 12 Y. Kohno, S. Saita, Y. Men, J. Yuan and H. Ohno, *Polym. Chem.*, 2015, **6**, 2163.
- 13 (a) H. Glasbrenner and H. Weingärtner, *J. Phys. Chem.*, 1989, **93**, 3378; (b) S. A. Buckingham, C. J. Garvey and G. G. Warr, *J. Phys. Chem.*, 1993, **97**, 10236; (c) H. Weingärtner, M. Kleemeier, S. Wiegand and W. Schröer, *J. Stat. Phys.*, 1995, **78**, 169; (d) M. Kleemeier, W. Schröer and H. Weingärtner, *J. Mol. Liq.*, 1997, **73–74**, 501; (e) Z.-L. Xie and A. Taubert, *ChemPhysChem*, 2011, **12**, 364; (f) Y. Kohno and H. Ohno, *Aust. J. Chem.*, 2012, **65**, 91; (g) Y. Kohno, Y. Deguchi and H. Ohno, *Chem. Commun.*, 2012, **48**, 11883; (h) S. Saita, Y. Kohno, N. Nakamura and H. Ohno, *Chem. Commun.*, 2013, **49**, 8988; (i) Y. Tsuji and H. Ohno, *Chem. Lett.*, 2013, **42**, 527; (j) T. Ando, Y. Kohno, N. Nakamura and H. Ohno, *Chem. Commun.*, 2013, **49**, 10248; (k) Y. Fukaya and H. Ohno, *Phys. Chem. Chem. Phys.*, 2013, **15**, 4066; (l) Y. Fukaya and H. Ohno, *Phys. Chem. Chem. Phys.*, 2013, **15**, 14941; (m) S. Saita, Y. Mieno, Y. Kohno and H. Ohno, *Chem. Commun.*, 2014, **50**, 15450; (n) Y. Deguchi, Y. Kohno and H. Ohno, *Chem. Lett.*, 2015, **44**, 238.
- 14 (a) J. E. L. Dullius, P. A. Z. Suarez, S. Einloft, R. F. de Souza and J. Dupont, *Organometallics*, 1998, **17**, 815; (b) P. J. Dyson, D. J. Ellis and T. Welton, *Can. J. Chem.*, 2001, **79**, 705; (c) C. A. Cerdeirinha, J. Troncoso, C. P. Ramos, L. Romani, V. Najdanovic-Visak, H. J. R. Guedes, J. M. S. S. Esperança, Z. P. Visak, M. Nunes da Ponte and L. P. N. Rebelo, Ionic Liquids III A, in *Fundamentals, Progress, Challenges, and Opportunities*, ed. R. Rogers and K. R. Seddon, ACS Symp. Ser., 2005, ch. 13, vol. 901, p. 175; (d) P. Nockemann, B. Thijs, S. Pittois, J. Thoen, C. Glorieux, K. V. Hecke, L. V. Meervelt, B. Kirchner and K. Binnemans, *J. Phys. Chem. B*, 2006, **110**, 20978; (e) Y. Fukaya, K. Sekikawa, K. Murata, N. Nakamura and H. Ohno, *Chem. Commun.*, 2007, 3089; (f) P. Nockemann, K. Binnemans, B. Thijs, T. N. Parac-Vogt,



K. Merz, A.-V. Mudring, P. C. Menon, R. N. Rajesh, G. Cordoyiannis, J. Thoen, J. Leys and C. Glorieux, *J. Phys. Chem. B*, 2009, **113**, 1429.

15 Y. Kohno and H. Ohno, *Phys. Chem. Chem. Phys.*, 2012, **14**, 5063.

16 U. Domańska, E. Bogel-Łukasik and R. Bogel-Łukasik, *Chem. – Eur. J.*, 2003, **9**, 3033.

17 L. Ropel, L. S. Belvèze, S. N. V. K. Aki, M. A. Stadtherr and J. F. Brennecke, *Green Chem.*, 2005, **7**, 83.

18 M. L. S. Batista, L. I. N. Tomé, C. M. S. S. Neves, E. M. Rocha, J. R. B. Gomes and J. A. P. Coutinho, *J. Phys. Chem. B*, 2012, **116**, 5985.

19 Y. Koga, *Solution Thermodynamics and Its Application to Aqueous Solutions: A Differential Approach*, Elsevier, Amsterdam, 2007, and references are cited in this textbook.

20 Y. Koga, *Phys. Chem. Chem. Phys.*, 2013, **15**, 14548.

21 Y. Koga, *J. Mol. Liq.*, 2015, **205**, 31.

22 P. Westh, H. Kato, K. Nishikawa and Y. Koga, *J. Phys. Chem. A*, 2006, **110**, 2072.

23 Y. Koga, P. Westh, K. Nishikawa and S. Subramanian, *J. Phys. Chem. B*, 2011, **115**, 2995.

24 J. Hu, W. M. Chiang, P. Westh, D. H. C. Chen, C. A. Haynes and Y. Koga, *Bull. Chem. Soc. Jpn.*, 2001, **74**, 809.

25 K. Miki, P. Westh and Y. Koga, *J. Phys. Chem. B*, 2005, **109**, 19536.

26 T. Morita, P. Westh, K. Nishikawa and Y. Koga, *J. Phys. Chem. B*, 2012, **116**, 7328.

27 K. Miki, P. Westh, K. Nishikawa and Y. Koga, *J. Phys. Chem. B*, 2005, **109**, 9014.

28 H. Kato, K. Nishikawa and Y. Koga, *J. Phys. Chem. B*, 2008, **112**, 2655.

29 H. Kato, K. Miki, T. Mukai, K. Nishikawa and Y. Koga, *J. Phys. Chem. B*, 2009, **113**, 14754.

30 T. Morita, A. Nitta, K. Nishikawa, P. Westh and Y. Koga, *J. Mol. Liq.*, 2014, **198**, 211.

31 E. Calvet and H. Prat, *Recent Progress in Microcalorimetry*, Pergamon Press, Oxford, 1963.

32 K. R. Loblich, *Thermochim. Acta*, 1994, **231**, 7.

33 B. Löwen, S. Schulz and J. Seippel, *Thermochim. Acta*, 1994, **235**, 147.

34 M. T. Parsons, P. Westh, J. V. Davies, C. Trandum, W. M. Chiang, E. G. M. Yee and Y. Koga, *J. Solution Chem.*, 2001, **30**, 1007.

35 I. Wadso and R. N. Goldberg, *Pure Appl. Chem.*, 2001, **73**, 1625.

36 T. Kondo, Y. Miyazaki, A. Inaba and Y. Koga, *J. Phys. Chem. B*, 2012, **116**, 3571.

37 Private communication.

38 T. Singh and A. Kumar, *J. Phys. Chem. B*, 2007, **111**, 7843.

39 J. Bowers, C. P. Butts, P. J. Martin, M. C. V. Gutierrez and R. K. Heenan, *Langmuir*, 2004, **20**, 2191.

40 J. Wang, H. Wang, S. Zhang, H. Zhang and Y. Zhao, *J. Phys. Chem. B*, 2007, **111**, 6181.

41 N. V. Sastry, N. M. Vaghela and V. K. Aswal, *Fluid Phase Equilib.*, 2012, **327**, 22.

42 L. Almásy, M. Turmine and A. Perera, *J. Phys. Chem. B*, 2008, **112**, 2382.

43 M. Tariq, F. Moscoso, F. J. Deive, A. Rodriguez, M. A. Sanromán, J. M. S. S. Esperança, J. N. C. Lopes and L. P. N. Rebelo, *J. Chem. Thermodyn.*, 2013, **59**, 43.

44 A. G. Avent, P. A. Chaloner, M. P. Day, K. R. Seddon and T. Welton, *J. Chem. Soc., Dalton Trans.*, 1994, 3405.

45 R. Wang, W. Leng, Y. Gao and L. Yu, *RSC Adv.*, 2014, **4**, 14055.

46 T. Morita, P. Westh, K. Nishikawa and Y. Koga, *J. Phys. Chem. B*, 2014, **118**, 8744.

47 S. Saita, Y. Kohno and H. Ohno, *Chem. Commun.*, 2013, **49**, 93.

48 Y. Koga, *J. Phys. Chem.*, 1996, **100**, 5172.

49 H. Katayanagi, K. Nishikawa, H. Shimozaki, K. Miki, P. Westh and Y. Koga, *J. Phys. Chem. B*, 2004, **108**, 19451.

50 C. M. Ellis, *J. Chem. Educ.*, 1967, **44**, 405.

51 M. D'Angelo, G. Onori and A. Santucci, *Chem. Phys. Lett.*, 1994, **220**, 59.

