



Cite this: *Phys. Chem. Chem. Phys.*,
2016, 18, 24377

Micelle formation of a non-ionic surfactant in non-aqueous molecular solvents and protic ionic liquids (PILs)[†]

Emmy C. Wijaya,^{ab} Frances Separovic,^a Calum J. Drummond^c and
Tamar L. Greaves^{*c}

Many ionic liquids and low molecular weight polar solvents have been reported to support amphiphile self-assembly, with most of these reported for the first time in the last decade. This phenomenon is attributed to the solvophobic effect (analogous to the hydrophobic effect in water). However, to date there has been no systematic study which evaluates micelle formation in a large library of non-aqueous solvents. Here we investigate micelle formation of a non-ionic amphiphile, hexa-ethyleneglycol mono *n*-dodecyl ether, C₁₂E₆, in a diverse range of molecular solvents and protic ionic liquids (PILs). Nine of the 19 non-aqueous molecular solvents investigated, and all four of the PILs, were found to support micelle formation. A link was investigated between the solvent cohesive energy density (as estimated using the Gordon parameter) and both the critical micelle concentration and the related free energy of micellization ($\Delta G_{\text{mic}}^{\circ}$). In addition, the chemical structure and liquid mesostructure of the solvent were found to be important factors in the ability of the solvents to support micelle formation.

Received 17th May 2016,
Accepted 8th August 2016

DOI: 10.1039/c6cp03332f

www.rsc.org/pccp

Introduction

The tendency of surfactants to aggregate in water, widely known as the hydrophobic effect, is significant for diverse applications such as enabling surfactants in soaps and detergent to remove oil-based stains and dirt. At a molecular level, the hydrophobic effect is a key factor in lipid bilayer formation in cell membranes, and maintaining the tertiary structure of proteins. The ability of solvents to promote amphiphile self-assembly is not limited to water, and over the last decades, there have been an increasing number of non-aqueous solvents which have been reported to show the analogous solvophobic effect. Previously we identified 16 commercially available, non-aqueous molecular solvents with solvophobic properties,¹ increasing to 24 the number of small molecular amines and glycols known to exhibit this property.^{2–4}

Ionic liquids (ILs), which are molten salts with melting points below 100 °C, are another non-aqueous solvent group where some have been reported to support amphiphile self-assembly.^{5–10} Some common features shared by many molecular solvents

and protic ILs (PILs) with a solvophobic effect are the presence of two or more potential hydrogen bonding centres, which enables them to form three-dimensional hydrogen bonded network structures. They usually have few, if any, methylene groups, and often have polar groups such as hydroxyls, both of which increases their solvophobic effect.³ While previously few solvents were considered amphiphile self-assembly media, it is now evident that many ILs, including PILs and aprotic ILs,¹¹ and molecular solvents share this property. It is anticipated that many more will be identified than are currently known.

One solvent property that is a useful indicator of the solvophobicity is the cohesive energy density.⁵ The cohesive energy density is the energy required to remove a unit volume of molecules completely from their neighbours to infinite separation (an ideal gas state). This is equal to the heat of vaporisation divided by the molar volume. For ILs in which it is not feasible to measure the heat of vaporisation due to negligible vapour pressures, the cohesive energy density can be approximated by taking into account the surface tension at the liquid–vapour interface, γ_{LV} , and the molar volume, V_{m} , as shown in eqn (1):

$$\text{Cohesive energy density} = c\gamma_{\text{LV}}/V_{\text{m}}^{1/3} \quad (1)$$

where c is a scalar quantity. The Gordon parameter, G , is obtained from eqn (1) when c is set to 1, as shown in eqn (2).^{6,12,13} G has been used to give a measure of the cohesive energy density, and

^a School of Chemistry, Bio21 Institute, University of Melbourne, VIC 3010, Australia

^b CSIRO Material Science and Engineering, Clayton VIC 3169, Australia

^c School of Science, College of Science, Engineering and Health, RMIT University,
GPO Box 2476, Melbourne VIC 3001, Australia.

E-mail: tamar.greaves@rmit.edu.au

[†] Electronic supplementary information (ESI) available. See DOI: 10.1039/c6cp03332f



thus the solvophobic effect for a variety of molecular and ionic liquids.¹⁴

$$G = \gamma_{LV}/V_m^{1/3} \quad (2)$$

Solvents with a high Gordon parameter generally have a higher solvophobic effect and hence are likely to be better at promoting amphiphile self-assembly.^{1,6,10,14} Water has the highest known G value of $2.743 \text{ J mol}^{1/3} \text{ m}^{-3}$, whereas other non-aqueous solvents have lower G values, with *N*-tert-butylformamide having the lowest at $0.53 \text{ J mol}^{1/3} \text{ m}^{-3}$ reported in the literature, while still retaining the ability to promote amphiphile self-assembly.⁵ Although not an absolute measure of the solvophobic effect between different non-supercritical solvents, and is a simple parameter that can be easily determined for most solvents. The strength of the solvophobic effect varies between solvents and hence the aggregation of amphiphiles within them varies, such as which lyotropic liquid crystal (LLC) mesophases are supported and what is the thermal stability range of the phases.¹

Studying micelle formation in solvents provides insight into the interactions between amphiphiles and solvents, and a comparison between solvents for their ability to support amphiphile self-assembly. An important micellisation parameter is the critical micelle concentration (CMC), which is defined as the surfactant concentration above which micelles spontaneously form.¹⁵ Currently, there is no systematic study that evaluates micelle formation in a large library of non-aqueous solvents. Instead the limited CMC data reported for surfactants in non-aqueous solvents is sporadic, and hence minimal comparisons can be made between solvents. However, from the data available it is clear that water is a more favourable media for micelle formation, shown by the smaller CMC values compared to those reported in non-aqueous solvents.^{2,3,8,16–20}

In this paper we have investigated the correlation between the solvophobic effect and micelle formation of a non-ionic amphiphile in a broad range of commercially available low molecular weight polar molecular solvents and in four PILs. These solvents were all selected because they have previously been shown to support LLC phases.¹ A protic ionic liquid, EAN, has been previously shown to support micelle formation.^{21,22} The chemical structure of the amphiphile hexa-ethyleneglycol mono *n*-dodecyl ether (C_{12}E_6) is provided in Fig. 1, and those of the molecular solvents and PILs provided in Fig. 2 and 3, respectively.

Experimental methods

Materials

Non-ionic surfactant, hexa-ethyleneglycol mono *n*-dodecyl ether, C_{12}E_6 (Nikkol BL-6SY) was supplied by Nikko Chemicals Co. Ltd,

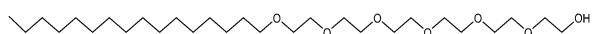


Fig. 1 Structure of the non-ionic surfactant, hexa-ethyleneglycol mono *n*-dodecyl ether (C_{12}E_6).

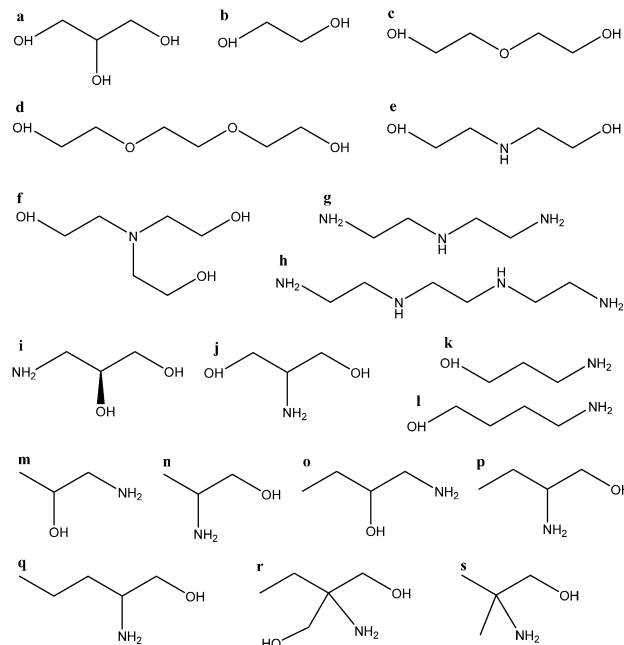


Fig. 2 Chemical structures of the molecular solvents: (a) glycerol, (b) ethylene glycol, (c) diethylene glycol, (d) triethylene glycol, (e) diethanolamine, (f) triethanolamine, (g) diethylene triamine, (h) triethylene tetramine, (i) 3-amino-1,2-propanediol, (j) 2-amino-1,3-propanediol, (k) 3-amino-1-propanol, (l) 4-amino-1-butanol, (m) 1-amino-2-propanol, (n) 2-amino-1-propanol, (o) 1-amino-2-butanol, (p) 2-amino-1-butanol, (q) 2-amino-1-pentanol, (r) 2-amino-2-ethyl-1,3-propanediol, and (s) 2-amino-2-methyl-1-propanol.

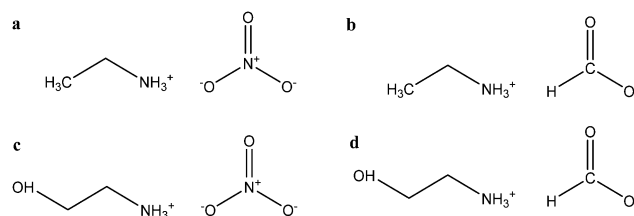


Fig. 3 Chemical structures of the PILs: (a) ethylammonium nitrate (EAN), (b) ethylammonium formate (EAF), (c) ethanolanionium nitrate (EOAN), and (d) ethanolanionium formate (EOAF).

Tokyo, Japan, at the highest purity. The 19 molecular solvents were used without further purification: glycerol (Merck, 99%), ethylene glycol (Sigma Aldrich, 99.8%), diethylene glycol (Sigma Aldrich, 99%), triethylene glycol (Sigma Aldrich, 99%) diethanolamine (AnalaR, 99.5%), and triethanolamine (Sigma Aldrich, 99.5%), diethylene triamine (Sigma Aldrich, 99%), triethylene tetramine (Sigma Aldrich, 97%), (*S*)-3-amino-1,2-propanediol (Sigma Aldrich, 97%), 2-amino-1,3-propanediol (Sigma Aldrich, 98%), 3-amino-1-propanol (Sigma Aldrich, 99+%), 4-amino-1-butanol (Sigma Aldrich, 98%), *DL*-1-amino-2-propanol (Sigma Aldrich, 90%), (*R*)-(-)-2-amino-1-propanol (Sigma Aldrich, 98%), 1-amino-2-butanol (Sigma Aldrich, 97%), 2-amino-1-butanol (Merck, 95%), *DL*-2-amino-1-pentanol (Sigma Aldrich, 97%), 2-amino-2-ethyl-1,3-propanediol (Acros Organics, 97%), and 2-amino-2-methyl-1-propanol (Fluka, 99%).



Ethylamine (Sigma-Aldrich, 70 wt%), ethanolamine (Chem Supply, 99.5%), nitric acid (Chem Supply, 70% w/w), and formic acid (Merck, 98–100%) were used without further purifications for synthesis of the PILs. Ethylammonium nitrate (EAN), ethylammonium formate (EAF), ethanolammonium nitrate (EOAN), and ethanolammonium formate (EOAF) were synthesized by dropwise addition of acid to the equimolar amount of amine at a temperature below 0 °C or above the freezing point of the reaction mixture during synthesis. Excess solvent was removed on a rotary evaporator, and samples were further dried using a benchtop freeze dryer (FreeZone).²³

The amount of water present in the molecular solvents and the synthesized PILs was measured using a Mettler Toledo DL39 Karl Fischer coulometer.

Experimental

Solvent–surfactant solutions were prepared using C₁₂E₆ in the various solvents over the concentration range of 0.0001 to 1 M. The solutions were mixed with a vortex mixer and left at room temperature for at least one hour before use. All solutions were examined using an optical microscope. Surfactant insolubility was indicated by the presence of “oil-like” droplets. Surface tension measurements were made of all these solutions using a pendant drop KSV Attension Tensiometer. 20 frames of 1 second intervals were acquired for each sample. These were repeated 5 times to ensure consistency. All measurements were performed at room temperature, 22 ± 1 °C.

Cloud-point measurements were made on solutions of 1 wt% C₁₂E₆ in each solvent heated to 100 °C, while visual observations were made at what temperature, if any, the solutions went cloudy.

Solutions of C₁₂E₆ in all molecular solvents and PILs for small angle scattering (SAXS) measurements were prepared from two times the CMC up to 10 M, with the upper limit depending on the amphiphile solubility in the different solvents. The solutions were then transferred into 1.5 mm special glass capillaries, and sealed with wax to avoid water absorption and solvent evaporation. Samples were analysed at the SAXS/WAXS beamline at the Australian Synchrotron, Melbourne, Australia. Spectra were acquired over the scattering vector range of 0.01 < *q* < 0.6 Å^{−1}, with 1 second exposures per sample. The scattering patterns were acquired at 25 °C, which is above the Krafft temperature of C₁₂E₆ of 20 °C in water.²⁴ The 2D SAXS scattering patterns were converted to 1D scattering curves with the SAXS I5-id software provided at the Australian Synchrotron. Background scattering curves were obtained from an empty capillary, which was subtracted from the sample scattering patterns. Software SasView²⁵ was used to fit the scattering curves in the *q* range of 0.03–0.3 Å^{−1} with different micelle models.

Micelle scattering models

The X-ray scattering intensity for micelles is given by eqn (3), where *I*(*q*) is the scattering intensity, *K* is a constant, *N* is the number of scatterers, *P*(*q*) is the form factor, and *S*(*q*) the structure factor.

$$I(q) = KNP(q)S(q) \quad (3)$$

The form factor *P*(*q*) and the structure factor *S*(*q*) describe the shape of the micelles and the interactions between micelles, respectively. Interactions between micelles are assumed to be insignificant for low micelle concentrations, *i.e.*, the structure factor is 1 and, thus, the scattered intensity is only described from the form factor. In relatively high concentrations, interactions between micelles needs to be taken into account; thus the structure factor needs to be included. Pedersen provides a comprehensive overview on SAXS scattering modelling that includes a variety of form factors and structure factors for different micelle shapes.²⁶

In this work we have used a variety of core–shell micelle models, including spherical, ellipsoidal, rod and disc, to fit the SAXS scattering data for C₁₂E₆ in the different solvents at a variety of concentrations. The models which gave the best fits, and the most logical sequence of parameters on changing concentration, were selected for each solvent–C₁₂E₆ combination. Micelle fitting in this work used a hard sphere structure factor when there was a high concentration of C₁₂E₆ present. This estimates the interparticle structure factor for monodisperse spherical particles which interact through excluded volume, using a Percus–Yevick closure,²⁷ and is the most suitable for the non-ionic amphiphile used in this work. In C₁₂E₆, the hydrocarbon core radius must not be greater than the fully extended C₁₂ chain length, which is 16.7 Å,²⁸ according to eqn (4), in which *n_c* is the number of carbons in the hydrocarbon chain.

$$l_{\max} = 1.5 + 1.265n_c \quad (4)$$

SAXS scattering originates from the electron density contrast between the micelle shell and core, and between the micelle shell and the solvent. The electron scattering length density (SLD) were calculated for the solvents using $SLD = \rho_e r_e$, where *r_e* is the electron radius, and $\rho_e = Z/V$, with *Z* the number of electrons and *V* the molecular volume. The SLD of the various solvents are provided in Table 3, and the calculation of the scattering length density for the core and shell of C₁₂E₆ micelles can be found in the ESI.† It was assumed that the hydrocarbon core of micelles is not solvated, thus the SLD of the core was fixed to 7.92 × 10^{−6} Å^{−2}. The solvents were expected to solvate the solvophilic headgroups of C₁₂E₆ and hence the electron density of the micelle shell will vary from those of just the headgroups. Contrast matching occurred to some extent for samples where the SLD of the solvent was comparable to the SLD of the micelle shells. For these samples simple (not core–shell) sphere, ellipsoid and cylinder models were found to fit best.

Results

CMC and micelle interfacial properties

The CMC of C₁₂E₆ in water and in nine of the molecular solvents was obtained using pendant drop surface tension measurements, and these are provided in Table 1 along with their interfacial properties. Similarly the values for C₁₂E₆ in the PILs are provided in Table 2. The initial surface tension values



Table 1 Gordon parameter, CMC, surface tension of neat solvent (γ_0), surface tension at CMC (γ_{cmc}), surface excess concentration (Γ_{max}), minimum area per molecule (A_{min}), standard free energy of micelle formation ($\Delta G_{\text{mic}}^\circ$), and standard free energy of adsorption ($\Delta G_{\text{ad}}^\circ$) for C_{12}E_6 in molecular solvents at room temperature

Solvents	Water content (wt%)	Gordon parameter ($\text{J mol}^{1/3} \text{ m}^{-3}$)	CMC (M)	γ_0 (mN m^{-1})	γ_{cmc} (mN m^{-1})	$\Gamma_{\text{max}} (\times 10^{-6} \text{ mol m}^{-2})$	$A_{\text{min}} (\text{\AA}^2)$	$\Delta G_{\text{mic}}^\circ (\text{kJ mol}^{-1})$	$\Delta G_{\text{ad}}^\circ (\text{kJ mol}^{-1})$
Water	100	2.743 ³⁸	7.93×10^{-5}	71.9 ³⁸	31.89	4.34	38	−23.0	-1.58×10^3
Ethylene glycol	0.65	1.356 ³⁹ 1.20 ⁴⁰	2.64×10^{-2}	48.0 ³⁹	31.34	2.34	71	−9.0	-1.19×10^3
3-Amino-1-propanol	1.70	1.07	1.56×10^{-1a}	45.3 44.7 ⁴¹	30.80	1.66	100	−4.6	-1.46×10^3
Diethylene glycol	0.95	0.98	1.82×10^{-1a}	44.4 55.10 ⁴¹	33.18	1.72	97	−4.2	-1.10×10^3
4-Amino-1-butanol	1.88	0.97	2.45×10^{-1a}	43.9 43.84 ⁴¹	34.20	1.88	88	−3.5	-0.88×10^3
Diethylene triamine	0.80	0.91	2.58×10^{-1a}	43.4 48.85 ⁴¹	33.89	2.19	76	−3.4	-0.73×10^3
Triethanolamine	0.14	0.91	8.07×10^{-2}	46.7 51.48 ⁴¹	32.24	2.35	71	−6.2	-1.05×10^3
(R)-(-)-2-Amino-1-propanol	0.15	0.90	1.87×10^{-1a}	38.3 47.03 ⁴¹	33.58	1.28	130	−4.2	-0.6×10^3
Triethylene glycol	0.52	0.88	1.44×10^{-1a}	45.1 45.13 ⁴¹	32.70	1.89	88	−4.8	-1.16×10^3
DL-1-Amino-2-propanol	0.45	0.86	4.69×10^{-1a}	36.7 36.5 ³ 48.68 ⁴¹	32.03	0.7	226	−1.9	-1.06×10^3

^a The Gibbs equation is derived on the basis of an assumption of a low amphiphile concentration. This assumption may no longer be valid when the CMC is greater than 0.1 M. However, the values for Γ_{max} , A_{min} , $\Delta G_{\text{mic}}^\circ$ and $\Delta G_{\text{ad}}^\circ$ have been included for all samples to enable relative comparisons between them.

of all these solvents, their Gordon parameters (G), and water contents are also provided in Tables 1 and 2. The values obtained in this experiment for the CMC of C_{12}E_6 in water and in EAN are 1.27×10^{-4} M and 4.46×10^{-2} M, which are in good agreement to what has been reported in the literature.^{8,29} The authors are not aware of any CMC measurement of this surfactant in any of the other solvents used in this work.

A representative plot of surface tension with respect to increasing C_{12}E_6 concentration in ethylene glycol is provided in Fig. 4, and the plots for all the other solvents are provided in Fig. S1–S14 of the ESI.† The CMC corresponds to the intersection of the two linear lines. Above this concentration, the surface tension value will no longer undergo significant change.

The pendant drop technique was not feasible for four of the molecular solvents where the surface tension was less than 34 mN m^{-1} , viz. 2-amino-1-butanol (32.2 mN m^{-1}), 2-amino-2-ethyl-1,3-propanediol (33.35 mN m^{-1}), 1-amino-2-butanol (31.1 mN m^{-1}), and 2-amino-1-pentanol (30.4 mN m^{-1}). Furthermore this technique was not applicable for the solvents with high viscosity, i.e., 3-amino-1,2-propanediol (4.3 Pa s) and glycerol (1.41 Pa s), or solids, 2-amino-2-methyl-1-propanol, 2-amino-1,3-propanediol and diethanolamine. An alternative technique of fluorescence, using pyrene to detect the presence of hydrophobic micelle core,³⁰ was trialled but due to fluorescence quenching it was not suitable for obtaining the CMC in these remaining solvents, particularly those containing amines.³¹

Table 2 Gordon parameter, CMC, surface tension of neat solvent (γ_0), surface tension at CMC (γ_{cmc}), surface excess concentration (Γ_{max}), minimum area per molecule (A_{min}), standard free energy of micelle formation ($\Delta G_{\text{mic}}^\circ$), and standard free energy of adsorption ($\Delta G_{\text{ad}}^\circ$) for C_{12}E_6 in protic ionic liquids at room temperature

Solvents	Residual water content (wt%)	Gordon parameter ($\text{J mol}^{1/3} \text{ m}^{-3}$)	CMC (M)	γ_0 (mN m^{-1})	γ_{cmc} (mN m^{-1})	$\Gamma_{\text{max}} (\times 10^{-6} \text{ mol m}^{-2})$	$A_{\text{min}} (\text{\AA}^2)$	$\Delta G_{\text{mic}}^\circ (\text{kJ mol}^{-1})$	$\Delta G_{\text{ad}}^\circ (\text{kJ mol}^{-1})$
Ethanolammonium formate (EOAF)	0.89	1.45	1×10^{-3a}						
Ethylammonium nitrate (EAN)	0.70	1.060 ¹⁰ 1.40 ⁴⁰	4.04×10^{-2}	47.3 ⁴²	35.7	1.41	117.65	−8.0	−141
Ethylammonium formate (EAF)	0.42	0.87	1.81×10^{-2}	42	30.8	3.00	55.41	−9.9	−594
Ethanolammonium nitrate (EOAN)	0.20	0.82	2.05×10^{-4}	65	28.3	11.4	11.19	−21.0	−432

^a Surface tension measurement was feasible for measuring the changes in surface tension of C_{12}E_6 in EOAF, but due to its high viscosity of 220 mN m^{-1} ,⁴² surface tension measurements using the pendant drop technique were technically difficult. However, from Fig. S13 in the ESI it can be determined that the CMC value would be around 1×10^{-3} M. Since the gradient could not be accurately obtained the interfacial properties could not be determined.



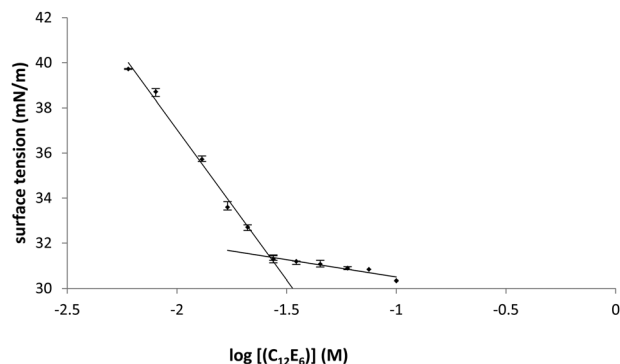


Fig. 4 Determination of CMC of $C_{12}E_6$ in ethylene glycol at room temperature using the pendant drop technique.

In addition, visual inspection by optical microscopy showed that $C_{12}E_6$ in triethylene tetramine formed an insoluble “oil-and-water”-like separation. Consequently the surface tension decrease with increasing concentration, shown in Fig. S10 (ESI[†]), was due to reaching a solubility limit rather than a CMC.

Surfactant insolubilities identification with cloud point measurement and optical microscopy

Cloud point measurements were undertaken in all the molecular solvents and PILs in the presence of 1 wt% $C_{12}E_6$. No cloudy appearance was observed in the molecular solvents up to 100 °C, except for water, in which the solution turned cloudy at 46 °C which was consistent with the literature value of 46.5 °C.³² Similarly there was no cloud point below 100 °C for EAF or EAN, consistent with the literature for EAN.³³ However, a cloud point at 1 wt% $C_{12}E_6$ in EOAN and EOAF was seen immediately at room temperature.

The absence of demixing (cloud point) below 100 °C for most of the non-aqueous solvents might be related to the solubility of the surfactant in these solvents. A previous study attributed the absence of a cloud point for $C_{12}E_5$, $C_{12}E_6$ and $C_{12}E_8$ in EAN below 130 °C to the hydrogen bonded network of EAN enhancing the solubility of the surfactant, and significantly increasing the cloud point temperature.³³ It is feasible that a similar phenomenon could occur for the non-aqueous solvents used in this investigation. Since they are solvents with polar moieties it is highly likely that there is an interaction with the ethoxy chain, increasing the solubility of surfactant molecules and increasing the cloud point temperature. In contrast, the cloud point seen in 1 wt% $C_{12}E_6$ in EOAF and EOAN at room temperature signifies reduced solubility of surfactant in these solvents, which might be due to the hydroxyl end group which could decrease the interaction between the alkyl chain of the surfactant and the solvents, making them more water like, and leading to the presence of a cloud point below 100 °C.

Micelle interfacial properties

The negative gradient of the surface tension with increasing concentration before the CMC is reached was used to provide insight into the micelle formation and packing of amphiphile molecules at the interface, according to the following equations.

The effectiveness of the surface tension reduction, Π_{cmc} , was calculated from eqn (5) for each solvent where the CMC was obtained using the surface tension technique.

$$\Pi_{cmc} = \gamma_0 - \gamma_{cmc} \quad (5)$$

where γ_0 is the surface tension of the neat solvent, and γ_{cmc} is the surface tension at the CMC. The minimum molecular area of the $C_{12}E_6$ surfactant adsorbed at the air–liquid interface, A_{min} , was calculated using a method based on the Gibbs adsorption equation.^{34–37}

$$A_{min} = 10^{20}/N_A \Gamma_{max} \quad (6)$$

where N_A is Avogadro's number and Γ_{max} is the maximum surface excess concentration at the air/liquid interface of the non-ionic surfactant, which takes into account the change in surface tension just before the CMC, described by eqn (7).

$$\Gamma_{max} = -1/2.303RT(d\gamma/d \log[C_{12}E_6]) \quad (7)$$

The standard free energy of micellization, ΔG_{mic}° , describes how energetically favourable the micellisation process is, thus giving a measure of the solvophobic interaction.

$$\Delta G_{mic}^\circ = -RT \ln(CMC) \quad (8)$$

The standard free energy of adsorption, ΔG_{ad}° , describes the free energy change for the adsorption of amphiphiles to the liquid–air interface, as in eqn (9). These values were calculated for each $C_{12}E_6$ –solvent system, and are shown in Tables 1 and 2. It should be noted that when $C_{12}E_6$ had a CMC greater than 0.1 M assumptions leading to the Gibbs equation may be challenged, but the values are considered to still provide useful indicative information.

$$\Delta G_{ad}^\circ = \Delta G_{mic}^\circ - (\gamma_0 - \gamma_{cmc})A_{min} \quad (9)$$

The values obtained for γ_{cmc} , A_{min} , ΔG_{mic}° , and ΔG_{ad}° of $C_{12}E_6$ in water and in EAN are in very good agreement with literature values.⁸ In all solvents, ΔG_{ad}° was calculated to be more negative compared to ΔG_{mic}° , which implies that the adsorption of amphiphiles at the interface is more spontaneous than micelle formation. Thus micelle formation starts after the completion of the adsorption phenomenon at the interface. Γ_{max} was positive in all solvents, which indicates that the concentration of $C_{12}E_6$ is higher at the liquid–air surface compared to in the bulk. This is typical behaviour of a surface active monomer, with the long hydrophobic chain preferring to be at the liquid–air interface compared to in the bulk solution.⁴³ The minimum molecular area, A_{min} , gives insight to the packing density of the surfactant in the interface.

For the molecular solvents, $C_{12}E_6$ had the smallest A_{min} of 38 Å² in water, hence the highest packing density, and corresponds to having the highest solvophobic interaction. The non-aqueous molecular solvents all had larger A_{min} values, and hence higher Γ_{max} values, which can be attributed to a weaker solvophobic interaction, which is directly proportional to ΔG_{mic}° . The larger A_{min} values in the non-aqueous solvents suggest the surfactant is aligning less perpendicular to the



liquid–air surface than in water, causing a less tightly packed surface. One possible explanation is that it is due to the reduction of solvent hydrophobicity at the surface, resulting in a drop of surface excess concentration,⁴⁴ which has previously been reported for binary mixtures of water and polar organic solvents.^{44,45}

$A_{\min}$ was the largest in 1-amino-2-propanol at $226 \text{ \AA}^2$, which suggests a much weaker solvophobic effect in this solvent, consistent with its low G parameter of $0.86 \text{ J mol}^{-1/3} \text{ m}^{-3}$. The large $A_{\min}$ is attributed to the methyl group in 1-amino-2-propanol interacting with the hydrophobic C_{12} alkyl chain of the surfactant.

The $A_{\min}$ for $C_{12}E_6$ in triethanolamine and ethylene glycol was $71 \text{ \AA}^2$. Despite its longer chain with an additional alkyl and hydroxyl branch, triethanolamine apparently does not incorporate into the surfactant headgroup. This might be due to its bulkier size leading to steric hindrance minimising its presence at the interface. Also, the terminal hydroxyl group will lead to decreased alkyl solubility, and drive it towards the bulk.

For the PILs, $\Gamma_{\max}$ was exceptionally high in EOAN compared to the other solvents. In fact the calculated $A_{\min}$ of $11 \text{ \AA}^2$ is significantly less than would be the case for a monolayer of molecules packed side by side with no intervening solvent and at 90° to the interface. Consequently the low $A_{\min}$ value indicates the adsorption of multilayers. $A_{\min}$ was higher in EAN compared to most solvents, except 2-amino-1-propanol. This might be due to the preferential orientation of EAN molecules in which the hydrocarbon moieties face the air.⁴² An interfacial monolayer microenvironment causes surfactants to orient so that the dodecyl chains are perpendicular to the interface and hence the ethylene oxide groups are more parallel to the interface in EAN compared to the other solvents.

Micelle identification by SAXS

Small and wide-angle X-ray scattering patterns (SAXS/WAXS) were acquired at 25°C and 50°C for each $C_{12}E_6$ –solvent combination for at least two concentrations above the CMC to confirm if micelles were present and, if so, determine their size and shape. The exceptions were $C_{12}E_6$ in 2-amino-2-methyl-1-propanol, 2-amino-1,3-propanediol, and diethanolamine due to these solvents being solids at room temperature.

A representative series of SAXS patterns is shown in Fig. 5 for $C_{12}E_6$ in ethylene glycol. The presence of micelles was evident by scattering at $q < 0.2 \text{ \AA}^{-1}$, which increased with increasing surfactant concentration. No scattering due to micelles was observable for $2\times$ CMC, weak scattering for $5\times$ CMC, and at 1 M the micelles are sufficiently concentrated for interactions to be observed, identified by the appearance of the characteristic peak at low q .

Among the 10 molecular solvents for which a CMC was obtained from pendant drop experiments there were only 6 that showed scattering consistent with micelles. In addition, 2-amino-2-ethyl-1,3-propanediol, one of the four molecular solvents which were not feasible for use with the pendant drop method, showed evidence of micelles. These solvents are listed in Table 3 and the scattering patterns were fitted with a variety

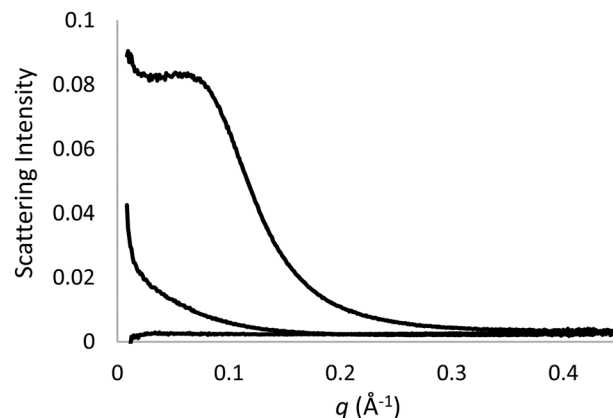


Fig. 5 SAXS patterns of ethylene glycol with $2\times$ CMC or 0.054 M (bottom), $5\times$ CMC or 0.135 M (middle), and 1 M $C_{12}E_6$ (top).

of micelle models, and the parameters giving the best fits and which were consistent across the different concentrations for each solvent, are provided in Table 3. The SAXS patterns for all $C_{12}E_6$ –solvent solutions are provided in Fig. S15–S21 of the ESI.†

The scattering patterns for $C_{12}E_6$ in all four PILs were consistent with the presence of micelles. However, for EOAF and EOAN, the scattering occurred at concentrations which were similar to the insolubility limit of the $C_{12}E_6$ in EOAF and EOAN, and consequently these spectra were not fitted. The parameters for the best fits to the micelles in EAN and EAF are provided in Table 3. The SAXS patterns for all $C_{12}E_6$ –PIL solutions are provided in Fig. S22 and S23 of the ESI.†

The scattering patterns all fitted well to spherical or elliptical micelles, either as simple or core–shell models, as can be seen in Table 3, with the models used for each concentration of $C_{12}E_6$ clearly specified. Micelles in ethylene glycol and triethanolamine fitted best to simple rather than core–shell models, which was due to similar electron scattering densities between the EO shell and these solvents. The micelles in all the other solvents fitted best to core–shell models. A hardsphere structure factor was included for the higher concentrations of $C_{12}E_6$ in many of the solvents to account for interactions between micelles. A volume fraction of 0.1 was obtained for those solvents in which hardsphere structure factor was used.

The overall radius of micelles in water, including the core radius and shell thickness, was $33.39 \text{ \AA}$, which is comparable to the previously reported value of $33 \text{ \AA}$.²⁸ Similar work performed by Penfold *et al.*, reported $5 \text{ wt\%}$ $C_{12}E_6$ formed micelles with a radius of $29 \text{ \AA}$,⁴⁶ with the smaller radius likely due to the inclusion of a polydispersity factor in their model. Of the non-aqueous molecular solvents, scattering data for micelles have only been reported in ethylene glycol, including small angle neutron scattering (SANS) on $10 \text{ wt\%}$ $C_{12}E_8$ in ethylene glycol, where Seguin *et al.* reported rod-shaped aggregates with a radius and length of 18 and $127 \text{ \AA}$, respectively.⁴⁷ These are comparable to our fits for $C_{12}E_6$ in ethylene glycol of oblate ellipsoids with an equatorial radius of $31\text{--}32 \text{ \AA}$, and polar radius of $7\text{--}8 \text{ \AA}$. The differences in radius and length are



Table 3 SAXS fitting parameters for C₁₂E₆ in molecular solvents and PILs. The form factor for the micelles was kept the same in each solvent

Solvent	[C ₁₂ E ₆] (M)	CMC (M)	Form factor	Structure factor	Polar radius (Å)	Equatorial radius (Å)	Shell thickness (polar)	Shell thickness (equatorial)	N _{agg}
Water	4.6 × 10 ⁻³	7.9 × 10 ⁻⁵	Coreshell spherical	—	14.3	—	19.6	—	35
	2.5 × 10 ⁻¹			Hardsphere	16.7	—	9.0	—	56
Ethylene glycol	1.4 × 10 ⁻¹	2.6 × 10 ⁻²	Ellipsoid	—	7.0	28.9			17
	1			Hardsphere	8.7	31.2			28
3-Amino-1-propanol	1	1.6 × 10 ⁻¹	Coreshell spherical	Hardsphere	12.8	—	8.6	—	25
Diethylene glycol	8.8 × 10 ⁻¹	1.8 × 10 ⁻¹	Coreshell spherical	Hardsphere	12.2	—	10.7		22
	1.5				13.4	—	7.6		28
Triethylene glycol	7.2 × 10 ⁻¹	1.4 × 10 ⁻¹	Coreshell ellipsoid	—	6.3	10.8	13.5	7.7	5
	1.5			Hardsphere	11.0	13.2	22.2	19.2	19
Triethanolamine	1	8.1 × 10 ⁻²	Ellipsoid	—	12.3	74.1	—		134
2-Amino-2-ethyl-1,3-propanediol	5	^a	Coreshell spherical	—	11.7	—	9.6		19
	10			Hardsphere	13.1	—	8.7		27
EAN	8.5 × 10 ⁻²	4.0 × 10 ⁻²	Coreshell spherical	—	9.3	—	3.6		10
	2.1 × 10 ⁻¹			Hardsphere	15.4	—	9.0		44
EAF	0.5	1.8 × 10 ⁻²	Coreshell ellipsoid	—	12.6	55.2	16.7	15.6	104

^a CMC unable to be obtained by surface tension measurement.

consistent with the larger ethoxy headgroup of their surfactant. In related work, Penfold *et al.*, fitted 5% C₁₂E₆ in various ethylene glycol–water solutions with a core–shell sphere of radius 16.7 Å and outer shell of 14 Å.⁴⁶ The use of a core–shell model was in contrast to our fits, and we attribute it to their only including ethylene glycol proportions up to 80 wt%, and hence we propose that for all their compositions there was sufficient water present to have a significant difference between the electron scattering density of the shell and the solvent.

All three EAF concentrations fit well to a coreshell oblate ellipsoid, with no structure factor required. The ellipsoidal micelles were observed to elongate with increasing C₁₂E₆ concentration, becoming more rod-like. The micelles in EAN fitted best to coreshell spheres, with a structure factor present at higher concentrations. Although the electron density between the EO shell and EAN are similar, models that did not include core–shell gave poorer fits. Similar work⁸ performed by Greaves *et al.* reported ellipsoidal micelles with an effective radius of around 12 Å at 10 wt% C₁₂E₆, which is comparable to the results in this work. Slight difference in micelle size might be due to the different shape model used. Employing the ellipsoidal model here for the 0.212 M C₁₂E₆ in EAN did not give a good fit.

Discussion

Here we have investigated the correlation between the solvophobic effect and micelle formation of a non-ionic amphiphile in a broad range of commercially available, low molecular weight polar molecular solvents and easily synthesised PILs. The non-ionic amphiphile, C₁₂E₆, was selected because it has a low Krafft temperature (20 °C) in water,²⁴ is soluble in a broad range of solvents, and has previously been reported to form micelles in water⁸ and EAN.^{14,48} This investigation enabled comparisons to be made between the non-aqueous solvents, their Gordon parameters and CMCs. We are not aware of any comparable investigation which has used such a broad range of solvents.

The CMC of C₁₂E₆ in all the non-aqueous solvents were at least one order of magnitude higher than in water, with the exception of EOAN, which had a CMC that was about 2.5 times larger than water. This reflects the weaker solvophobic interaction in non-aqueous solutions compared to the hydrophobic effect in aqueous solutions. In general the CMCs in the PILs were lower than in the non-aqueous molecular solvents. It was apparent from the lower Gibb's free energy of micelle formation, $\Delta G_{\text{mic}}^{\circ}$, in non-aqueous solvents compared to water that there was a lower tendency for micellar aggregation. This can be attributed in part to the nonpolar interactions between the alkyl chains of C₁₂E₆ and nonpolar moieties of the solvents. These interactions increase the solubility of C₁₂E₆ and consequently lead to higher CMC values. The rough correlation between the Gibb's free energy of micellisation and the solvophobic effect within the molecular solvents and PILs was shown by plotting $\Delta G_{\text{mic}}^{\circ}$ against the Gordon parameter, *G* (Fig. 6). It is apparent for the molecular solvents that there is a roughly monotonic relationship, inferring that more energy is generally needed for micelle formation in solvents with lower Gordon parameter values. It is also very clear from Fig. 6 that water has a substantially higher Gordon parameter, and hence higher solvophobic effect, compared to the other solvents used in this work, together with a correspondingly favourable free energy of micellization. The four PILs all had more favourable $\Delta G_{\text{mic}}^{\circ}$ values compared to the molecular solvents with the same Gordon parameters. This suggests that the interaction between cations and anions in PILs are stronger than the interactions in the solvent structures of molecular solvents. Furthermore, the CMC of C₁₂E₆ in EOAF and EOAN of $\sim 1 \times 10^{-3}$ and 2.05×10^{-4} M, respectively, was significantly smaller than in EAF and EAN of 4.25×10^{-2} and 7.81×10^{-2} M, respectively. We attribute this difference to the effect of the hydroxyl group in EOAN and EOAF, which reduced the solubility of C₁₂E₆ in these PILs and hence leads to lower CMC values.

The specific chemical groups present in the molecular solvents and PILs had a large impact on whether or not they



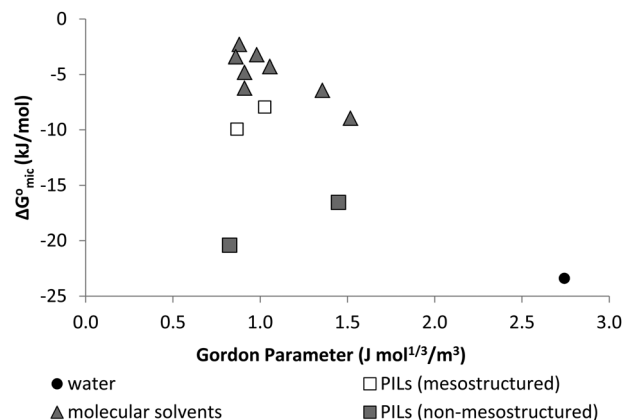


Fig. 6 Gordon parameter versus $\Delta G^\circ_{\text{mic}}$ of C_{12}E_6 in the molecular solvents and PILs.

supported the formation of C_{12}E_6 micelles, and if so, on the CMC. Overall, solvents with hydroxyl groups were better solvents for micelle formation compared to those with amine groups, and increasing the proportion of hydrocarbons, particularly methyl or ethyl groups, was highly detrimental. For example, the $\Delta G^\circ_{\text{mic}}$ of C_{12}E_6 in hydroxyl containing diethylene glycol and amine containing diethylene triamine was -4.22 and -3.36 kJ mol^{-1} , respectively, showing the higher driving force for micelle formation in the hydroxyl containing solvent, even though they had similar G (0.98 $\text{J mol}^{1/3} \text{ m}^{-3}$ for the former and 0.91 $\text{J mol}^{1/3} \text{ m}^{-3}$ for the latter).

Addition of a methylene group increased $\Delta G^\circ_{\text{mic}}$ from -4.60 kJ mol^{-1} in 3-amino-1-propanol to -3.48 kJ mol^{-1} in 4-amino-1-butanol, due to the increased solubility of the amphiphile. This trend was also seen with $\Delta G^\circ_{\text{mic}}$ going from -9.00 to -4.22 kJ mol^{-1} from ethylene glycol to diethylene glycol with the increased proportion of methylene. The $\Delta G^\circ_{\text{mic}}$ of C_{12}E_6 in triethylene glycol did not continue this trend, with its value of -4.81 kJ mol^{-1} lying between those of ethylene glycol and diethylene glycol. We suggest this may be due to steric hindrance preventing the alkyl chains of the amphiphile to interact with the hydrocarbon portion of the solvent.

The effect of increasing the alkyl chain length on the solvent had a more dramatic effect in reducing the solvent ability to support amphiphile self-assembly compared to increased methylene groups with terminal polar groups. This could most clearly be observed since no molecular solvent which had a terminal ethyl group supported amphiphile self-assembly. For example, the methyl containing 1-amino-2-propanol and 2-amino-1-propanol both supported micelles, whereas their ethyl and propyl counterparts of 1-amino-2-butanol, 2-amino-1-butanol and 2-amino-1-pentanol did not. The exception was 2-amino-2-ethyl-1,3-propanediol where scattering from micelles was detected using SAXS. The solvents 1-amino-2-butanol, 2-amino-1-butanol and 2-amino-1-pentanol have previously been shown to form a mesostructure consisting of polar and non-polar domains, whereas 2-amino-2-ethyl-1,3-propanediol did not.¹ Consequently it is likely that the non-polar domains within the mesostructured solvents may enhance the solubility

of amphiphiles and decrease the likelihood of micelle formation. In 2-amino-2-ethyl-1,3-propanediol the two hydroxyl and one amine group are likely to form a multiple hydrogen bonded network with steric hindrance making the segregation of the alkyl chains energetically unfavourable. Interestingly, contrary to the trend found in molecular solvents, EAN and EAF, which contain ethyl groups, were found to support micelle formation. This further shows the exceptional properties that PILs have compared to the molecular solvents.

The driving force for micelle formation of C_{12}E_6 in molecular solvents was significantly weaker than in the PILs. This was shown by the higher $\Delta G^\circ_{\text{mic}}$ in molecular solvents. Since the nonionic surfactant C_{12}E_6 will not be affected by the ionic behavior of the PILs, we suggest this is due to that stronger solvent-solvent interactions in PILs compared to in molecular solvents. Interestingly, the magnitude of A_{min} in these molecular solvents was found to be within the same range as the A_{min} in EAN and EAF (between 50 to 120 $\text{\AA}^2$), indicating that these molecular solvents and two PILs also behave as co-surfactants, and are incorporated to some extent within the micelle.

The effect of solvent mesostructure was also clearly observable for the four PILs. The hydroxyl containing EOAF and EOAN are not mesostructured, whereas EAN and EAF are ref. 49 and 50. The formation of non-polar domains would result in some solvation of the hydrocarbon of the amphiphiles and decrease the driving force for micelles to form, which is evident from the higher $\Delta G^\circ_{\text{mic}}$ in EAF and EAN compared to EOAN and EOAF.⁵¹ In addition, the larger A_{min} in EAF and EAN suggests that the alkyl chains of the cation behave to some extent as co-surfactants. In contrast, the non-structured EOAN, has a significantly higher Γ_{max} and lower A_{min} compared to EAF, EAN, water and the other molecular solvents. The particularly low A_{min} value of 11 $\text{\AA}^2$ indicates that the C_{12}E_6 formed multilayers at the air-liquid interface prior to micellization. It is presumed that similar amphiphile behavior would be present in EOAF.

The presence of micelles detected by SAXS can be related to solvent solvophobicity. In most cases, micelle features were only observed at concentration > 0.1 M for the non-aqueous solvents, which is higher than a concentration equivalent to $5 \times \text{CMC}$. The clearest example was seen in 2-amino-2-ethyl-1,3-propanediol in which micelles were only observed at a C_{12}E_6 concentration of 5 M, and the characteristic peak due to inter-micellar interactions was observed at 10 M. The shell thickness gives insight into the interaction between the EO shell and solvents, and was found to be the highest in water compared to either molecular solvents or PILs. This might be due to water being a better solvent for the EO headgroups, allowing them to extend more freely. It may also be an artefact due to the significantly higher contrast between the electron scattering length density of the solvent and the shell. Alternatively, the EO chains could be more compact in the other solvents, be it in the molecular solvents or PILs. This would indicate less interaction between the EO shell with the solvents, as found by Atkin *et al.*, in EAN.⁵²

The aggregation number (N_{agg}) describes the number of surfactant molecules in micelles once the CMC has been



reached. In general N_{agg} was observed to increase as the surfactant concentration increased (Table 3). Exceptions were micelle sizes in ethylene glycol and 2-amino-2-ethyl-1,3-propanediol in which micelle dimensions, and consequently N_{agg} , underwent a slight reduction with increasing surfactant concentration. This shows that in some cases micelle size does not depend on surfactant concentration. We consider that the micelles may have reached their maximum size due to the high surfactant concentrations used in this study. Consequently the effect of solvent solvophobicity and solvent chemical constituents on micelle size was difficult to determine from the limited data obtained and the variety of CMC values in these solvents. The solvents that can be compared are ethylene glycol, diethylene glycol and triethylene glycol in which the data of micelle size and N_{agg} are available at $5\times$ CMC. For these three solvents, N_{agg} decreased in the order $\text{EG} > \text{DG} > \text{TG}$, which corresponds to a decrease in solvent solvophobicity. In the case of the PILs, data for micelles at $2\times$ and $5\times$ CMC in EAN and EAF were obtained. From Table 3, micelles in EAF, which had lower G (and thus lower solvent solvophobicity) than EAN, had smaller micelle size and lower N_{agg} value. This also suggests that PILs with higher solvophobicities result in larger micelle sizes and N_{agg} values.

Overall, as indicated by lower CMC and $\Delta G_{\text{mic}}^\circ$ values, PILs used in this investigation were better at supporting micelle formation than the non-aqueous molecular solvents. Moreover, micelles were detected in EAN and EAF at a surfactant concentration as low as $2\times$ CMC, in which the size was comparable to micelles formed in 1 wt% in water. Interestingly, EAN, EAF, EOAF and EOAN had Gordon parameter values in the same range as for the molecular solvents. The ionic interaction between positive and negative ions in PILs are likely to be much stronger compared to interactions between molecular solvent, which would translate to lower CMC values, $\Delta G_{\text{mic}}^\circ$ and smaller micelle sizes.

Conclusions

The solvophobic effect and its effect on amphiphile self-assembly were explored by studying the ability of several non-aqueous polar organic solvents to support micelle formation. Generally, an increase in solvophobicity (indicated by higher G) resulted in lower CMC, and consequently lower $\Delta G_{\text{mic}}^\circ$ values. Calculation of interfacial properties suggest that, compared to water, the selected non-aqueous solvents cause surfactant molecules to align less perpendicular to the air–water interface, as indicated by larger A_{min} , or may form multilayers, as indicated by significantly smaller A_{min} values. Micelle size and N_{agg} were found to decrease as G decreased. This shows that solvent solvophobicity is a significant contributor to micelle formation and that G is a useful parameter for predicting the probability of amphiphile self-assembly in a solvent. However, there are other factors that need to be considered, such as solvent mesostructure and specific chemical moieties. This study also shows that, based on lower CMCs, select PILs are better solvents

for promoting amphiphile self-assembly compared to the molecular solvents. This indicates that the ionic interaction between cations and anions in PILs are stronger when compared to the specific hydrogen-bond interactions between molecular solvents.

Acknowledgements

ECW is the recipient of a University of Melbourne International Fee Remission Award, a CSIRO Materials Science and Engineering PhD Studentship and Albert Shimmins Writing Up Award. This research was undertaken in part on the SAXS/WAXS beamline at the Australian Synchrotron, Melbourne, Australia.

Notes and references

- 1 E. C. Wijaya, T. L. Greaves and C. J. Drummond, *Faraday Discuss.*, 2013, **167**, 191.
- 2 A. Ray, *J. Am. Chem. Soc.*, 1969, **91**, 6511.
- 3 A. Ray, *Nature*, 1971, **231**, 313.
- 4 T. L. Greaves, A. Weerawardena and C. J. Drummond, *Phys. Chem. Chem. Phys.*, 2011, **13**, 9180.
- 5 T. L. Greaves and C. J. Drummond, *Chem. Soc. Rev.*, 2013, **42**, 1096.
- 6 T. L. Greaves, A. Weerawardena, C. Fong and C. J. Drummond, *J. Phys. Chem. B*, 2007, 4082.
- 7 T. L. Greaves and C. J. Drummond, *Chem. Soc. Rev.*, 2008, **37**, 1709.
- 8 T. L. Greaves, S. T. Mudie and C. J. Drummond, *Phys. Chem. Chem. Phys.*, 2011, **13**, 20441.
- 9 T. L. Greaves, A. Weerawardena, I. Krodkiewska and C. J. Drummond, *J. Phys. Chem. B*, 2008, **112**, 896.
- 10 T. L. Greaves, A. Weerawardena, C. Fong and C. J. Drummond, *Langmuir*, 2007, **23**, 402.
- 11 I. A. Sedov and B. N. Solomonov, *Fluid Phase Equilib.*, 2016, **425**, 9.
- 12 D. Kabir ud and P. A. Koya, *Langmuir*, 2010, **26**, 7905.
- 13 P. Ajmal Koya, D. Kabir ud and K. Ismail, *J. Solution Chem.*, 2012, **41**, 1271.
- 14 D. F. Evans, *Langmuir*, 1988, **4**, 3.
- 15 D. Yu, F. Huang and H. Xu, *Anal. Methods*, 2012, **4**, 47.
- 16 H. N. Singh, S. M. Saleem, R. P. Singh and K. S. Birdi, *J. Phys. Chem.*, 1980, **84**, 2191.
- 17 M. S. Akhter and S. M. Alawi, *Colloids Surf., A*, 2003, **219**, 281.
- 18 A. Lattes and I. Rico, *Colloids Surf.*, 1989, **35**, 221.
- 19 M. S. Akhter, *Colloids Surf., A*, 1999, **150**, 25.
- 20 M. S. Akhter, S. M. Alawi and A. N. Bose, *Colloids Surf., A*, 1995, 173.
- 21 D. F. Evans, A. Yamauchi, R. Roman and E. Z. Casassa, *J. Colloid Interface Sci.*, 1982, **88**, 89.
- 22 D. F. Evans, A. Yamauchi, G. J. Wei and V. A. Bloomfield, *J. Phys. Chem.*, 1983, **87**, 3537.
- 23 T. L. Greaves, A. Weerawardena, I. Krodkiewska and C. J. Drummond, *J. Phys. Chem. B*, 2008, **112**, 896.
- 24 T. Inoue, Y. Higuchi and T. Misono, *J. Colloid Interface Sci.*, 2009, **338**, 308.



- 25 SAS Institute; SAS: The Power to Know 2016, http://www.sas.com/en_au/home.html, date accessed 5 May 2016.
- 26 J. S. Pedersen, *Adv. Colloid Interface Sci.*, 1997, **70**, 171.
- 27 J. K. Percus and G. J. Yevick, *Phys. Rev.*, 1958, **110**, 1.
- 28 C. Tanford, *The Hydrophobic Effect: Formation of Micelles and Biological Membranes*, Wiley, New York, 1973.
- 29 R. Dong and J. Hao, *Chem. Rev.*, 2010, **110**, 4978.
- 30 J. Aguiar, P. Carpena, J. A. Molina-Bolívar and C. Carnero Ruiz, *J. Colloid Interface Sci.*, 2003, **258**, 116.
- 31 B. K. Paul and S. P. Moulik, *Ionic liquid-based surfactant science: formulation, characterization and applications*, John Wiley & Sons, Hoboken, New Jersey, 2015.
- 32 K. S. Sharma, S. R. Patil and A. K. Rakshit, *Colloids Surf., A*, 2003, **219**, 67.
- 33 M. U. Araos and G. G. Warr, *J. Phys. Chem. B*, 2005, **109**, 14275.
- 34 C. J. Drummond, G. G. Warr, F. Grieser, B. W. Ninham and D. F. Evans, *J. Phys. Chem.*, 1985, **89**, 2103.
- 35 E. Matijevic and B. A. Pethica, *Trans. Faraday Soc.*, 1958, **54**, 1382.
- 36 M. J. Rosen and S. B. Sulthana, *J. Colloid Interface Sci.*, 2001, **239**, 528.
- 37 C. Fong, T. L. Greaves, T. W. Healy and C. J. Drummond, *J. Colloid Interface Sci.*, 2015, **449**, 364.
- 38 D. J. Mitchell, G. J. T. Tiddy, L. Waring, T. Bostock and M. P. McDonald, *J. Chem. Soc., Faraday Trans. 1*, 1983, **79**, 975.
- 39 T. Wolff and G. von Büнау, *Ber. Bunsen-Ges. Phys. Chem.*, 1984, **88**, 1098.
- 40 A. H. Beesley, D. F. Evans and R. G. Laughlin, *J. Phys. Chem.*, 1988, **92**, 791.
- 41 C. L. Yaws, *Thermophysical Properties of Chemicals and Hydrocarbons*, William Andrew Inc., Norwich, N.Y., 2008.
- 42 T. L. Greaves, A. Weerawardena, C. Fong, I. Krodziewska and C. J. Drummond, *J. Phys. Chem. B*, 2006, **110**, 22479.
- 43 I. Mukherjee, S. P. Moulik and A. K. Rakshit, *J. Colloid Interface Sci.*, 2013, **394**, 329.
- 44 S. Das, S. Mondal and S. Ghosh, *J. Chem. Eng. Data*, 2013, **58**, 2586–2595.
- 45 S. Sinha, D. Tikariha, J. Lakra, A. Tiwari, S. Saha and K. Ghosh, *J. Surfactants Deterg.*, 2015, **18**, 629.
- 46 J. Penfold, E. Staples, I. Tucker and P. Cummins, *J. Colloid Interface Sci.*, 1997, **185**, 424.
- 47 C. Seguin, J. Eastoe, R. Clapperton, R. K. Heenan and I. Grillo, *Colloids Surf., A*, 2006, **282**, 134.
- 48 M. U. Araos and G. G. Warr, *Langmuir*, 2008, **24**, 9354.
- 49 R. Atkin and G. G. Warr, *J. Phys. Chem. B*, 2008, **112**, 4164.
- 50 T. L. Greaves, D. F. Kennedy, N. Kirby and C. J. Drummond, *Phys. Chem. Chem. Phys.*, 2011, **13**, 13501.
- 51 T. L. Greaves, D. F. Kennedy, S. T. Mudie and C. J. Drummond, *J. Phys. Chem. B*, 2010, **114**, 10022.
- 52 O. Werzer, G. G. Warr and R. Atkin, *Langmuir*, 2011, **27**, 3541.

