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Dispersions of Magnetic Nanoparticles in the Mixture Ethyleneglycol-Choline Chloride : the Role of Solvent Association

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Abstract

The mixture of choline chloride and ethyleneglycol ((1:3) on the molar scale) is first characterized using density, viscosimetry and refractive index measurements, in the temperature range 20-45°C, with controlled amounts of water. Thermodynamic parameters such as the activation entropy, enthalpy and Gibbs energy for the viscous flow are deduced from the data and show that the solvent is highly structured. A protocol to disperse in this solvent 10 nm maghemite nanoparticles previously synthesized in water is described. The good quality of the dispersion is assessed by visual observations, optical microscopy, small angle X-ray scattering (SAXS), magnetization measurements, viscosimetry, and light scattering. The results lead to the description of a local viscosity around the particles, lower than the macroscopic one, that could be the result of the solvent segregation with ethyleneglycol being mostly present in the close vicinity of the particles.

Keywords:

Magnetic nanoparticles
Colloidal dispersions
Choline chloride
Ethyleneglycol
Maghemite

Introduction

Dispersions of nanoparticles (NPs) in ionic liquids (ILs) have drawn recently huge attention. The electrochemical and conducting properties of the ILs can indeed be combined with specific (electronic, magnetic, optical...) properties of the nanoparticles to obtain multiple stimuli responsive materials with possible applications in electrochemistry, synthesis, catalysis, separation fields [1, 2].

However, one serious drawback of such solvents is their price, toxicity and availability. For this reason, deep eutectic solvents (DES) and related complex solvents can be interesting alternatives. These liquids are made of a quaternary ammonium salt (choline chloride in our case) mixed with a molecular hydrogen bond donor (HBD). Since the molar ratio of ammonium salt in the final liquid is huge, the new solvent is closer to an ionic liquid than to an electrolyte solution. Indeed, the physico-chemical properties of DESs such as negligible vapor pressure, thermostability, wide electrochemical potential windows, tunability are similar to those of the traditionally used ionic liquids. DESs can also have the ILs' drawbacks such as high viscosity [3]. Since the interest for DESs is more recent than that for standard ILs, data about both the DESs and dispersions of particles in DESs are still scarce [4, 5]. The complexity of the solvent,

made of three components (the cation, the anion and the HBD) makes any prediction about colloidal stability in DESs impossible in the actual state of our knowledge. For this reason and due to the interesting properties of DESs such as cost and availability there is a huge need for experimental data about dispersions of NPs in DESs.

In this work, we use a solvent mixture made of choline chloride (ChCl), a widely used, cheap, bio-sourced component, mixed with ethyleneglycol (EG) as HBD. Thanks to its composition, it is closely related to DES and is sometimes classified among them although it is not an eutectic mixture on the strict basis of melting temperatures. This solvent will be named ChEG hereafter. Though somehow toxic, EG is chosen since it can form with ChCl a liquid of low viscosity at room temperature (around 35 mPa.s at 20°C for ChCl:EG (1:3 in mol:mol) compared to more than 1000 mPa.s for the widely used ChCl:urea (1:2)[6]). Based on the analysis of the conductivity and viscosity, it was shown [7] that ChCl:EG mixtures switch from a dilute salt solution behavior to an ionic liquid behavior for ChCl:EG composition about 1:4 (20 mol% = 36 vol% of ChCl). For this reason, a composition slightly higher (1:3 - 25 mol% = 43 vol% of ChCl) is used in this work.

A dispersion of NPs in ChEG could be obtained by direct synthesis in the complex solvent. This leads to NPs with surface properties different from the synthesis in water and to enhanced catalytic properties [8, 9]. However, the nature of the surface

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is not very well understood and controlled. Moreover, it can be difficult to eliminate the side-products of the NPs synthesized in ChEG.

Here, we chose a NPs synthesis in an aqueous medium which is easier, cheaper and already well known. Therefore, a transfer to ChEG after a synthesis in water appears more advantageous, all the more so since the interface NPs-solvent can be controlled and adapted during this step. The chosen nanoparticles are iron oxide maghemite particles (diameter ≈ 10 nm), synthesized by coprecipitation in alkaline aqueous medium [10, 11, 12]. Their surface state can be tuned to monitor the surface charge by changing the pH, or by adsorption of various ligands such as citrate or polyacrylate. Such particles have already been successfully dispersed in an ionic liquid, ethylammonium nitrate [11, 13, 14, 15]. The particles being magnetic, the observed stable dispersion is indeed a ferrofluid (FF). The best colloidal stability was obtained for particles coated with citrate or poly(acrylate), using Na^+ as a counterion [11, 15]. The stability was ensured by the formation of layers of solvent ions around the particles, providing a steric hindrance to particles approach [14]. The determining effect of small amounts of water was also evidenced, leading to enhanced flocculation by deconstruction of these protective layers [14].

In the present work, we transfer similar citrate-coated NPs (Na^+ counterions) from water to ChEG. In the first part, we detail the samples and techniques used. We then analyse the solvent - mixture ChCl-EG (1:3) - by measuring viscosity, density and refractive index in the temperature range 20-45°C, as a function of the water content. These parameters are indeed necessary for the analysis of the obtained colloidal dispersions in ChEG. A model to describe the results is proposed. This solvent is subsequently used to disperse maghemite nanoparticles. We assess the colloidal stability by means of visual observations, optical microscopy, small angle X-ray scattering (SAXS), magnetization measurements, viscosimetry, and dynamic light scattering. Finally, we discuss the results and conclude on the possible influence of a local viscosity different from the macroscopic one to understand the data.

1. Materials and methods

The following products are purchased and used as received: choline chloride (Sigma Aldrich); ethyleneglycol (Normapur, VWR); nitric acid (69.5% water solution, Carlo Erba); hydrochloric acid (37% water solution, AnalaR Normapur, VWR); citric acid (H_3Cit , Rectapur, Prolabo); iron(II) chloride (AnalaR Normapur, VWR); iron(III) chloride (Prolabo); iron(III) nitrate (Technical, VWR); trisodium citrate (Na_3Cit , Merck); acetone (technical, VWR); diethyl ether (99.8% pur., AnalaR Normapur).

Water content is analyzed using coulometric Karl Fisher titration (Schott TitroLine KF Trace). Density is measured thanks to an Anton Paar DSA 5000M, with precision 10^{-6} g/mL and temperature accuracy better than 10^{-3} °C. Viscosity of ChEG and ChEG/water mixtures is determined using an Anton Paar Automated MicroViscometer (AMVn) instrument by using the falling ball model in a 1.8 mm (resp. 3.0 mm)

capillary with a 1.5 mm (resp. 2.5 mm) falling ball. The instrument is first calibrated with reference standard oils N14 (resp. N44) from Cannon Instrument Company provided by Anton Paar ($\eta = 71.29$ mPa·s (resp. 19.68 mPa·s), $\rho = 0.8249$ g/mL (resp. 0.8092 g/mL) at 25°C)). Dynamic light scattering is performed on a Vasco instrument from Cordouan Technologies, operating at $\lambda = 656$ nm, with a temperature control better than 0.1°C. The instrument uses a very thin film (around 200 μm) and a backscattering detection at $\theta = 135^\circ$, which avoids multiple scattering and allows measurements in optically absorbing samples such as ferrofluids. The corresponding scattering vectors Q values are $Q = 2.3 \cdot 10^{-3} \text{ \AA}^{-1}$ in water and $2.5 \cdot 10^{-3} \text{ \AA}^{-1}$ in ChEG. The index of refraction is measured with a refractometer Arago from Cordouan Technologies at 656 nm, with a temperature control better than 0.1°C. The samples are observed on the micron scale with an optical microscope (Olympus BX51 with lens 40x/0.6 and camera x5). Small angle X-rays scattering (SAXS) is performed at the SWING beam line of synchrotron Soleil (France). Two different configurations are used, giving access to the Q -range $2.10^{-3} \text{ \AA}^{-1} < Q < 0.4 \text{ \AA}^{-1}$. The beam energy of 7 keV (corresponding to $\lambda = 1.77 \text{ \AA}$) is chosen to avoid X-ray absorption by iron. Standard correction procedures are applied for sample volume, empty cell signal subtraction and detector efficiency to obtain the scattered intensity in absolute scale (cm^{-1}). Such data reduction is done using the software Foxtrot[®]. Magnetization is registered using a home-made Vibrating Sample Magnetizer (VSM [16, 17]). The stability under magnetic field is checked from the absence of diffraction pattern when the sample is illuminated by a He-Ne laser under the field of an electromagnet. Above the threshold of stability, a diffracted line appears in the direction perpendicular to the magnetic field due to the formation of concentrated phase needles resulting from a phase separation [18].

2. Choline Chloride-Ethyleneglycol (ChEG)

The complex solvent chosen here, based on choline chloride (ChCl) and ethyleneglycol (EG) is prepared by simple mixture of the two components at 60°C in closed bottles in order to avoid water uptake. No liquid mixture could be observed for a molar ratio ChCl:EG of composition (1:1)(50mol%) even at high temperature and crystals appeared upon cooling for the ratio (1:2)(33mol%). For this reason and according to Abbott's criterion to obtain an ionic liquid behaviour [7], we chose to work with a molar ratio (1:3). The properties of both components and resulting solvent are given in Table 1. The amount of water in the prepared solvent, determined from Karl Fisher titration, is 0.2 w%.

Only few data are available in literature (see Table 1), with scattered values. Our density is close to the results of [7]. However, the viscosity we obtain is much higher. Since the water contents can be of importance for the viscosity, we investigate the density and viscosity of the ChEG, in the temperature range 20-45°C (Figure 1 and 2, data in SI) for various quantities of

Properties	Ethylene glycol (EG)	Choline Chloride (ChCl)	ChCl:EG 1:3 (ChEG)
M (g/mol)	62.07	139.63	81.46
Melting T (°C)	-13 [19]	247 [20]	-8 [21]
Boiling T (°C)	197.3 [19]	decomposition	
Density (g/cm ³)	1.10988 [22] 1.11004 [23]	1.1 [20] (for 70w% in water)	1.1367 [21] 1.118 [7] 1.115167*
Viscosity (10 ⁻³ Pa/s)	17.13 [24]	-	26 [21] 19 [7] 31.6*
Refractive index	1.4318 (20°C) [19]	-	1.456*
Molar volume (cm ³ /mol)	55.92 [22]	127 ± 5 [20]	71.66 [21] 72.86 [7] 72.53*

Table 1: Properties of the individual components of ChEG and of ChEG at T=25°C except when another temperature is specified. * = this work

added water. Unless a huge amount of water is present in literature samples, this cannot be the reason for the observed viscosity discrepancies¹.

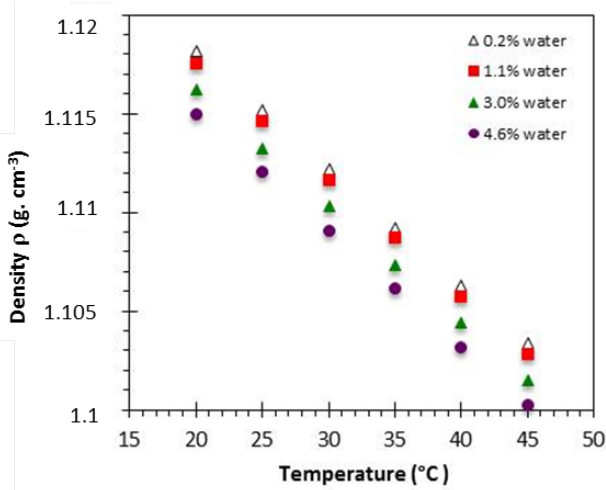


Figure 1: Density versus temperature for the solvent ChCl:EG (1:3) with different water contents (in weight percent).

The evolution of the viscosity with the water content of Fig. 2 shows the importance of controlling this parameter. Indeed these solvents are hygroscopic. We left a vial open to the ambient air and recorded the mass increase, attributed to atmospheric water uptake, for several days. The water uptake is very low (+0.24 w%) during the first four hours. It is thus possible to work quickly with short opening of the containers for the necessary handling of ChEG. In the long run, the water uptake is +13 w% after 3 weeks. Thus the solvent must be

¹Another reason for the low viscosity values found in the literature could be a degradation of the solvent, for example due to a heating at too high temperature, as this solvent presents a thermal decomposition above 100°C [25]

stored in appropriate conditions to avoid water contamination. The subsequent elimination of water by freeze-drying for instance is not possible, since EG is eliminated as well.

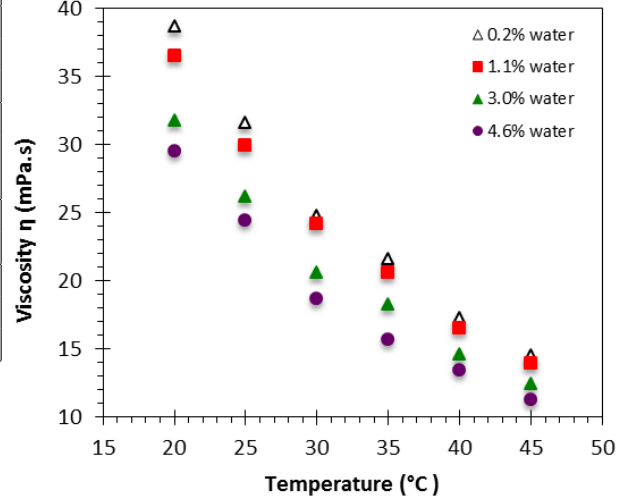


Figure 2: Viscosity versus temperature for the solvent ChCl:EG (1:3) with different water contents (in weight percent).

2.1. Dependence of the density on the water-content as a function of temperature

Molar volume of the solvent (here, wet ChEG) V_{solv} can be calculated from experimental density values ρ for each temperature according to :

$$V_{solv} = M_{solv} / \rho \quad (1)$$

(M_{solv} is the molar mass of the solvent)

The quantity of water is taken into account in M_{solv} through:

$$M_{solv} = (1 - x)M_{ChEG} + xM_{H_2O} \quad (2)$$

where x is the molar fraction of water deduced from Karl-Fisher titration, M_{ChEG} and M_{H_2O} are, respectively, the molar mass of the pure ChEG (table 1) and water. V_{solv} vs water content x is plotted in supporting information (Fig. SI-1), for temperatures between $T=20^\circ\text{C}$ and 45°C . The plots are linear and can be approximated for each T by :

$$V_{solv} = V_{ChEG}^* + x(V_{H_2O}^\infty - V_{ChEG}^*) \quad (3)$$

where $V_{H_2O}^\infty$ is the partial molar volume of water at infinite dilution and V_{ChEG}^* is the molar volume of pure (dry) ChEG, at the given temperature T . The y-intercept is thus V_{ChEG}^* at T . From the slopes of the curves, one can calculate $V_{H_2O}^\infty$ as a function of T . Table 2 reports these V_{ChEG}^* and $V_{H_2O}^\infty$ as well as that of the molar volume of pure water $V_{H_2O}^*$ calculated from [26], in the temperature range explored. $V_{H_2O}^\infty$ is smaller by 1 cm³ compared to $V_{H_2O}^*$ at 20°C and stays small whatever the temperature.

The same decrease was observed for water dissolved in pure EG [27] and can be attributed to a more efficient packing of water in the mixture with less H bonds around H₂O that in bulk water.

T	20	25	30	35	40	45
V_{ChEG}^*	72.840	73.035	73.230	73.425	73.620	73.817
$V_{H_2O}^\infty$	16.994	17.083	17.174	17.266	17.361	17.487
$V_{H_2O}^*$	18.032	18.053	18.079	18.108	18.141	18.178

Table 2: Molar volume of pure ChEG V_{ChEG}^* (cm³/mol), partial molar volume of water at infinite dilution in the ChEG-water mixture $V_{H_2O}^\infty$, and molar volume of pure water $V_{H_2O}^*$ for various temperature T (°C). Assuming an ideal mixture, V_{ChEG}^* is consistent with a molar volume of 72.0±1.5 cm³/mol at 20°C which can be estimated from the pure component volumes with an imprecise density of choline chloride.

The expansion of V_{slvt} with temperature can be quantified with the volumetric isobaric thermal expansion coefficient, defined as $\alpha_P = (1/V_{slvt})(\partial V_{slvt}/\partial T)_P$. The coefficient α_P values are between $5.27 \cdot 10^{-4}$ and $5.36 \cdot 10^{-4} \text{ K}^{-1}$ in the temperature range explored whatever the water content. These values are close to those determined for the mixture ChCl/EG (1:2) ($5\text{--}6 \cdot 10^{-4} \text{ K}^{-1}$) [28] and a bit smaller than the one of EG ($6.4 \cdot 10^{-4} \text{ K}^{-1}$) [24]). This indicates that ChEG is a slightly more associated liquid than pure EG.

2.2. Dependence of viscosity on the water-content as a function of temperature

The temperature-dependence of macroscopic viscosities η of ChCl:EG (1:3) mixtures (Fig. 2) can be described with Eyring's model [29] for each water content as:

$$\eta = hN_A/V_{slvt} \exp(\Delta G_a/RT) \quad (4)$$

with h , the Planck constant, N_A , the Avogadro number and ΔG_a the activation Gibbs energy for the viscous flow. From equation 4, ΔG_a is obtained for each x and T using the experimental η and V_{slvt} .

In a first step, let us analyse ΔG_a at 298 K as a function of the water molar fraction x . We obtain a linear function, which can be described with $\Delta G_a = -22,9x + 21,5$ (kJ/mol). The y-intercept (21.5 kJ/mol) corresponds to ΔG_a value for pure (dry) ChEG. The presence of water lowers the activation Gibbs energy for the flow, i.e., the energy barrier to overcome for the fluid to flow is smaller. This reduction of ΔG_a can originate either from a smaller enthalpy contribution (intermolecular interactions in the fluid are smaller) or from an entropic contribution (more configurations are accessible) during the flow process according to Eyring's hole theory for viscosity.

In a second step, the analysis of ΔG_a as a function of T for each water content gives the activation flow enthalpy and entropy from $\Delta G_a = \Delta H_a - T\Delta S_a$. When a molecule is on its way to change location (producing a flow), it must overcome an energy barrier made of two contributions. First, it must fight against the loss of interaction energy with its neighbours of the initial place : this is quantified as ΔH_a . Second, since it is not linked any more to its neighbours, it gains rotation entropy. This

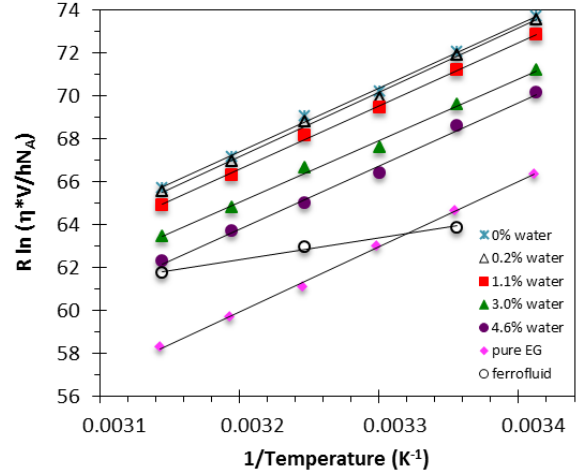


Figure 3: Eyring's plot for the solvent ChCl:EG (1:3) with different water contents (in weight percent). 0% was obtained using extrapolated values of molar volumes and viscosities to null water content. Values for pure EG are from [24]. For the ferrofluid, the molar volume is that of the driest ChEG (0.2% water) with the local viscosity discussed in the text.

is ΔS_a . The flow will be easier if its activation energy ΔG_a is low, e.g., if ΔH_a and $-T\Delta S_a$ are low (high ΔS_a). These contributions can be determined according to equation (4) supposing ΔH_a and ΔS_a are independent of T. $R \ln[\eta V_{slvt}/hN_A] = \Delta G_a/T$ plotted vs $(1/T)$ (Eyring plot, figure 3) is a straight line which gives access to ΔH_a (slope) and ΔS_a (y-intercept) (Table 3). This is applied for each water content as well as for the extrapolated "pure dry solvent" using V_{ChEG}^* and viscosity data linearly extrapolated to zero water content (data in SI). All ΔH_a values are scattered around the mean value 29.5 ± 0.5 kJ/mol, close to the enthalpy of activation of pure EG (30.0 kJ/mol)[24]. This implies that, for the viscous flow, the formation of an activated species in the solvent made of ChCl and EG is probably related to the same kind of intermolecular forces as in pure EG. The presence of some water does not deeply modify this interaction energy. It is coherent with a view of the cohesion of ChEG due to H bonds between all three components (EG, ChCl and water).

ΔS_a values are scattered around $28.5 \pm 1.5 \text{ J mol}^{-1} \text{ K}^{-1}$ (or $-T\Delta S_a = -8.5 \pm 0.6 \text{ kJ mol}^{-1}$ at 298K) for all water contents. The entropic contribution $-T\Delta S_a$ to ΔG_a is negative and is about one third of ΔH_a as it is also observed for pure EG ($-T\Delta S_a = -10.7 \text{ kJ mol}^{-1}$ at 298K [24]). The entropy term is high and lowers the ΔG_a value. However, the solvent, i.e. the mixture of EG and ChCl, is still a structured solvent that needs to locally destructurate in order to flow, as pure EG. A more detailed analysis of the contributions of $-T\Delta S_a$ and ΔH_a to ΔG_a with the water content in ChEG is delicate since a small experimental imprecision on the slope determination in Figure 3 (i.e., ΔH_a) has a huge impact on the y-intercept ΔS_a .

This set of data on the ChCl:EG (1:3) mixture provides a good picture of this new solvent, scarcely described in the literature, here studied within several conditions of water uptake. Its stability in the long run for $T < 30^\circ\text{C}$ was also verified by

	ChEG (1:3) *	Pure EG [24]	FF in ChEG *
ΔG_a (kJ.mol ⁻¹) (298K)	21.4	19.3	19
ΔH_a (kJ.mol ⁻¹)	29.9	30.0	10.0
$-T\Delta S_a$ (kJ.mol ⁻¹) (298K)	-8.5	-10.7	9.0

Table 3: Flow activation Gibbs energy, enthalpy and entropy derived from viscosity for the ChEG solvent with 0.2% water, for EG and for the ferrofluid in ChEG. * this work

regular checking of the samples over several months.

3. Dispersions in ChEG

The dispersions are produced by transfer of the NPs from water, solvent used for their synthesis, to ChEG. When a black liquid is obtained as seen by eye, it has to be analyzed further to evaluate the quality of the dispersion and its nanostructure. Micrometric structure is analyzed by optical microscopy. If the dispersion is homogeneous at this scale, small-angle X-ray scattering (SAXS), magnetization measurements and dynamic light scattering (DLS) are used.

3.1. Preparation of the dispersions

Iron oxide nanoparticles are synthesized by a routine process of coprecipitation of ferric and ferrous iron ions in alkaline aqueous medium [10, 11, 12]. It produces magnetite (Fe₃O₄) nanoparticles then fully oxidized to maghemite (γ-Fe₂O₃) which is chemically stable. Final dispersed nanoparticles with a diameter of the order of 10 nm are obtained in acidic aqueous medium (pH = 1.5). At this stage, the surface charge is positive (Fe-OH₂⁺) and counterbalanced by NO₃⁻ anions. The volume fraction of the NPs in the aqueous stock solution is Φ_v = 2.25v%, as deduced from the determination of the iron content by Flame Atomic Absorption Spectroscopy and using a density of 5g/cm³ for maghemite. Before their transfer to ChEG, the surface of the particles is covered by citrate ions, using the procedure described in [30]. Briefly, citric acid (H₃Cit) is added to the acidic dispersion which leads to flocculation. HNO₃ is then removed by repeated additions/removals of H₃Cit at 0.020 mol/L. The pH is then increased between 7.0 and 7.5 by addition of NaOH. The final particles are thus covered with citrate ions, Na⁺ being the positively charged counterions. This dispersion of citrate-coated NPs will be referred to as "aqueous dispersion" from now on.

If the particles are to be transferred in the ChEG, the aqueous dispersion is first flocculated by addition of acetone, washed 3 times by mixtures of acetone:water (9:1) and once by ether. After the removal of the ether, the desired amount of ChEG is added, and the dispersion is gently heated to remove any residual organic solvent. The dispersion is then centrifuged 10 minutes at 4000 rpm to get rid of the remaining few undispersed particles.

3.2. Structure by optical microscopy and Small Angle X-Ray Scattering (SAXS)

Two samples are compared, in water and in ChEG. As the aqueous systems have been extensively studied in several conditions and their colloidal stability is well controlled (see for example [31] and references therein), the dispersion in water is used as a reference. On the nanoscale, the structure is determined by SAXS. The scattered intensity *I* is recorded as a function of the scattering vector *Q*. For very dilute NPs dispersion, *I*(*Q*) is characteristic of the shape of the scattering objects over the whole *Q* range (form factor). For more concentrated dispersions, the nanoparticles undergo colloidal interactions. The intensity at small *Q*s is then representative of the structure factor *S*(*Q*) of the dispersion while at high *Q*s, it is still dominated by the form factor. A dilute aqueous sample is used for the determination of the form factor. It is obtained by dilution of the initial aqueous dispersion in HNO₃ 10⁻² mol L⁻¹. Figure SI-2 plots the scattered intensity versus the scattering vector *Q* in absolute units. Figure 4 shows the corresponding Porod plot *I*×*Q*⁴, normalized by the volume fraction Φ_v and the contrast Δρ², versus *Q*. The volume fraction is taken from the invariant of diffusion : Φ_v is 0.061 vol% for the reference sample in water and 1.13 vol% for the sample in ChEG. This representation *I*×*Q*⁴ enhances the high *Q*s region and allows the comparison of the form factors. The curves in water and in ChEG are very similar, indicating that the same particles are present in both solvents.

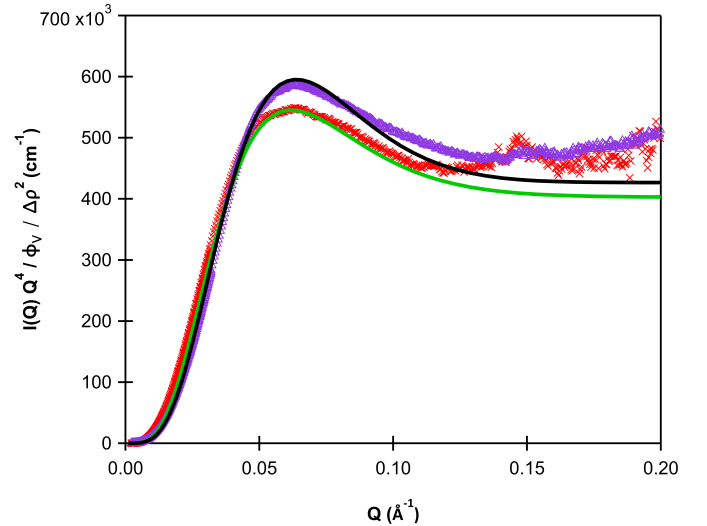


Figure 4: Porod (*I*×*Q*⁴) representation of SAXS data, in absolute unit. (red crosses) Bare particles in aqueous HNO₃ (pH = 2) : Φ_v = 0.061v% (purple triangles) Citrate-coated particles in ChEG : Φ_v = 1.13v%. Fits were obtained with a lognormal distribution for the sizes of the particles. See text for details.

At such a low concentration of NPs in water, we can assume that the interparticle interactions are negligible. We adjust the curve *I*(*Q*) with a form factor of polydisperse spheres and a log-normal distribution. The scattering length densities ρ necessary to calculate the contrast Δρ² are known (NPs : 40.6 10¹⁰cm⁻²;

water $9.3 \cdot 10^{10} \text{cm}^{-2}$; ChEG $10.2 \cdot 10^{10} \text{cm}^{-2}$) therefore the only parameters are the median diameter d_{SAXS} and the polydispersity σ_{SAXS} . We use the software provided by the NIST² to model the experimental curves [32]. The fits are performed in the $I \times Q^4$ representation (Figure 4). Good fits are obtained in water for $d_{SAXS} = 6.0 \text{ nm}$, $\sigma = 0.43$ and in ChEG for $d_{SAXS} = 6.0 \text{ nm}$, $\sigma = 0.40$. These results confirm that the size distribution is little affected by the solvent transfer. The quality of the fit is poorer at small Q values, indicating the presence of some aggregates in both cases (Figure SI-2).

The colloidal stability in ChEG means that some interparticular repulsions are present to counterbalance the van der Waals attraction. Information can be gained on the interactions from the scattered intensity at small Qs. The structure factor of the ChEG dispersion is calculated using the diffractogram in water (i.e., at very low concentration) as form factor : $S(Q) = [I_{ChEG}(Q)/\phi_{v,ChEG}]/[I_{water}(Q)/\phi_{v,water}]$. Indeed, we know from previous studies that interparticular interactions in water are negligible at this ϕ [33]. It is reported on figure 5. It presents an initial value $S(0)$ lower than 1 and no structure peak at larger Qs, which is indicative of short range repulsions. The order of magnitude of the repulsion is similar to what was observed with the same kind of particles (maghemite covered with citrate, Na^+ counterions) dispersed in the ionic liquid ethylammonium nitrate [11].

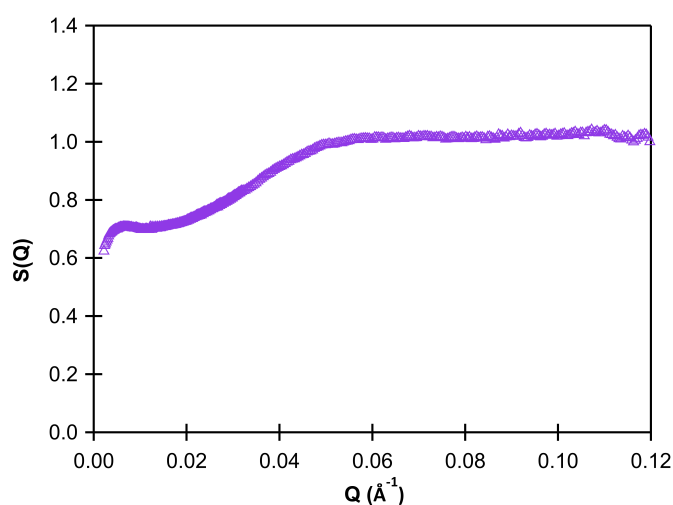


Figure 5: Structure factor $S(Q)$ of the dispersion in ChEG at $\Phi_v = 1.13\%$, using the diffractogram in water as form factor

3.3. Magnetization measurements

As the NPs are magnetic, the analysis of the magnetization curve of the dispersions can give information on the size distribution. However, the stability under field is checked first: both samples are stable up to $1350 \text{ Oe} = 108 \text{ kA m}^{-1}$ at room

temperature. Above this threshold, the destabilization occurs in a few ten minutes. Therefore, the magnetization versus magnetic field is determined using a sequence in high fields as quick as possible in order not to destabilize the system during the measurement. The curves, plotted in Figure 6, can be fitted with the Langevin's equation coupled with a lognormal distribution of NP diameters [34]. We get here 314 kA/m for the magnetization of the material, $d_o = 7.1 \text{ nm}$ and $\sigma = 0.38$ for the sample in water and $d_o = 6.8 \text{ nm}$ and $\sigma = 0.35$ for the sample in ChEG. This distribution in ChEG is thus very close to the initial distribution in water. It means that only few large particles are removed during the transfer in ChEG, as was already concluded from the SAXS measurements.

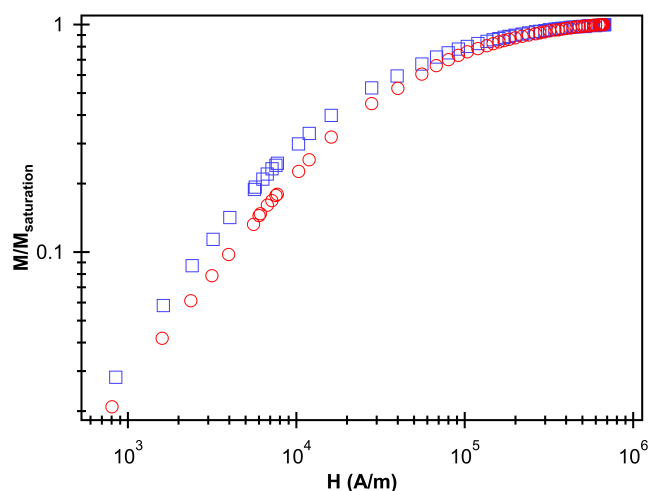


Figure 6: Normalized magnetization versus magnetic field for the samples in water (squares) and ChEG (circles).

3.4. Dynamic Light Scattering (DLS)

Much more accessible than SAXS, DLS is a laboratory experiment which enables studies with temperature, as well as aging and reversibility exploration. The light scattered by the sample is recorded versus time and an autocorrelation function is calculated. It measures how fast a NPs forgets its initial position thus giving access to a diffusion coefficient, which can be interpreted using appropriate models. Here, the measured intensity autocorrelation function $g_2(t)$ is converted into the field autocorrelation $g_1(t)$ function using : $g_1(t) = [g_2(t)]^{1/2}$ - offset. The function $g_1(t)$ of the studied samples can be fitted with a stretched exponential $g_1(t) = A \times \exp[-(t/\tau)^\beta]$ and an exponent β close to 1, however not perfectly, especially at long times. Henceforward, we choose (i) to fit the curves for times corresponding to $g_1(t) > 0.1$, without taking into account the tail which is less reliable, (ii) to use a mono-exponential, i.e., $\beta = 1$, in order to reduce the number of fitting parameters to only one. This corresponds to the simplest analysis, with a monodisperse population. The characteristic time τ is related to the translational diffusion coefficient $D_t = (\tau Q^2)^{-1}$.

²NIST : National Institute of Standards and Technology - https://www.ncnr.nist.gov/programs/sans/data/red_anal.html

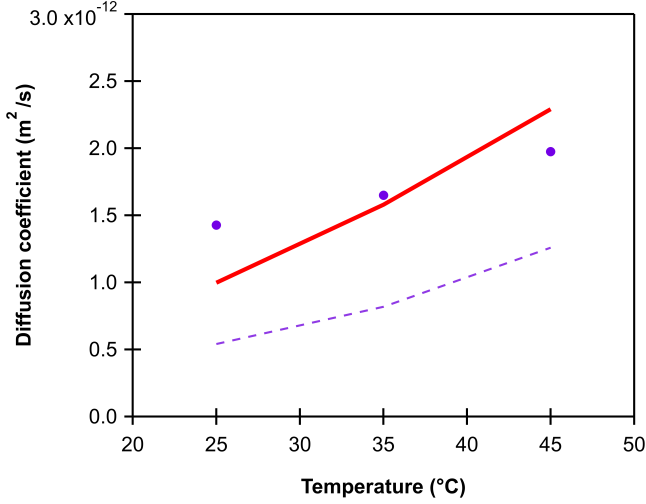


Figure 7: Diffusion coefficients D_t from DLS measurements. Points : experimental values $D_{t,ChEG}^{exp}$ in ChEG at various T. Dashed line : calculated values $D_{t,ChEG}^{calc}$ using the measured viscosities of ChEG. Solid line : calculated values $D_{t,EG}^{calc}$ using the viscosity of pure EG .

The DLS measurements in ChEG are performed on the same sample as SAXS at different temperatures , 25°C, 35°C and 45°C, for which macroscopic ChEG viscosities are known. As shown in Table 4 (and Figure 7), the diffusion coefficient D_t increases with temperature, in a reproducible manner, even for 6 months old samples for which no evolution is seen, and going back to its value at 25°C after a cycle 25°C - 45°C - 25°C.

Solvent	Water	ChEG	ChEG	ChEG
T (°C)	25	25	35	45
$D_{t,exp}$ ($m^2 s^{-1}$)	1.92 10^{-11}	1.43 10^{-12}	1.65 10^{-12}	1.97 10^{-12}
η_{macro} (mPa.s)	0.89	31.6	21.6	14.5
d_H^{app} (nm)	25.5	9.6	12.4	16.8
η_{local} from DLS (mPa.s)		12	10.7	9.2

Table 4: Experimental diffusion coefficients $D_{t,exp}$, macroscopic viscosities η_{macro} , corresponding hydrodynamic diameter d_H . Viscosities η_{local} are determined from DLS for the samples with the hypothesis that $d_H = 25.5$ nm for all T in ChEG.

4. Discussion

4.1. Hydrodynamic NP diameter and local viscosity in the ChEG-FF sample

When interparticle interaction is negligible, the translational diffusion coefficient D_t is related to the hydrodynamic diameter d_H by the Stokes Einstein's equation

$$D_t = \frac{k_B T}{3\pi\eta_0 d_H} \quad (5)$$

where k_B is the Boltzmann constant, T the absolute temperature, η_0 the local solvent shear viscosity.

Since D_t is obtained by DLS as an intensity average, the corresponding hydrodynamic diameter obtained through equation 5 is a mean value defined as $d_H = \langle d^6 \rangle / \langle d^5 \rangle$ [35]. It can be compared to the value calculated from the lognormal distribution obtained from SAXS, d_H^{calc} . In the case of the aqueous FF-sample, using for η_0 the macroscopic viscosity of water in Eq. 5, we obtain $d_H^{calc} = 17.0$ nm in reasonable agreement with the DLS experimental results $d_H = 25$ nm (see Table 4). The DLS value is indeed somehow higher due to the presence of small aggregates (see SAXS section).

However in the case of the ChEG FF-sample, this agreement fails if η_0 is assimilated to the ChEG macroscopic viscosity in Eq. 5. The apparent diameter d_H^{app} is then found much smaller than d_H^{calc} . Moreover, the size is unexpectedly temperature dependent.

Several arguments can be put forward to explain these observations. (i) A much lower polydispersity in the ChEG-FF sample could be an explanation, however not consistent with the SAXS results. (ii) Interparticle interactions could modify the apparent diffusion coefficient in the ChEG FF-sample, however the observed size decrease would correspond to a strong repulsion which is not compatible with SAXS data. (iii) A bad evaluation of the refractive index could be put forward, however the refractive indexes of liquids usually range roughly between 1.3 and 1.5 and their variation cannot explain such a large difference. (iv) The viscosity is not correct. Indeed, viscosity highly depends on the solvent composition. However a macroscopic water pollution cannot explain the results as a huge amount of water (more than 20%) would be necessary to induce such a large decrease of the macroscopic viscosity.

The only possibility in this ChEG-FF sample, is thus a lower local viscosity around the NPs than the macroscopic ChEG viscosity. A direct comparison of the diffusion coefficients is presented in Figure 7. It plots the temperature-dependence of the experimental diffusion coefficients $D_{t,exp}^{ChEG}$ and compares them with calculated values, using the sample in water at 25°C as a reference : $D_{t,calc}^{ChEG} = D_{t,exp}^{H_2O}(25) \times \eta_{macro}^{H_2O}(25) / \eta_{macro}^{ChEG}(T)$. The same can be done to estimate the diffusion coefficient of the particles in pure EG : $D_{t,calc}^{EG} = D_{t,exp}^{H_2O}(25) \times \eta_{macro}^{H_2O}(25) / \eta_{macro}^{EG}(T)$. Figure 7 shows that neither the macroscopic value of the viscosity of ChEG nor that of EG can reproduce the experimental data. A local viscosity is thus necessary around the particles to assess for the diffusion results. It can be determined through $\eta_{local}^{ChEG}(T) = D_{t,exp}^{H_2O}(25) / D_{t,exp}^{ChEG}(T) \times \eta_{macro}^{H_2O}(25)$ (see Table 4). We observe that this local viscosity is much lower than the macroscopic one and presents a less pronounced temperature dependence.

4.2. Analysis of the temperature-dependence and origin of the local viscosity in the ChEG-FF sample

The analysis with the model of Eyring of the temperature dependence of the local viscosity, considering that the molar volume of the solvent is unaffected by the presence of the particles, gives contributions to the flow Gibbs energy very different from those of pure ChEG (Figure 3 and Table 3). If at

298K the Gibbs energy value ΔG_a is similar in the ChEG-FF and in the complex solvent (around 19 kJ/mol), its splitting into enthalpy ΔH_a and entropy ΔS_a is completely modified. Indeed, ΔH_a is lowered from 30 to 10 kJ/mol while $-T\Delta S_a$ changed sign from -8.5 kJ/mol to +9 kJ/mol. This implies that the particles deeply modify the solvent organization around them, both from the solvent-solvent interaction point of view (a decrease of ΔH_a means that a lower number of interactions between solvents molecules must be modified for the fluid to flow) and from the entropic point of view (higher $-T\Delta S_a$ means that the movement of a solvent entity implies a loss of accessible configurations).

Ionic liquids are known to be strongly organized in hydrophilic and hydrophobic domains [36] and the DES made of choline chloride-urea was recently shown to be highly structured through H-bonds implying all its three components (Cl-, Cholinium, urea) [37]. By analogy with ChCl-urea, ChCl-EG can be assumed structured. In such a structured solvent, the nanoparticles can preferentially accommodate one of the solvent components and disturb the network. In this case, the viscosity experienced by the particles is not the macroscopic viscosity of the solvent, but the local viscosity of the surrounding fluid in that domain. In our case, this local viscosity is much lower than the macroscopic one, our results suggesting that the less viscous fluid, ethyleneglycol, directly surrounds the particles.

5. Conclusion

The density and viscosity of a mixture of choline chloride and ethylene glycol (1:3) a complex solvent close to DESs, were carefully determined from 20 to 45°C and for water contents between 0.2 and 4.6%. The analysis of the data using the model of Eyring indicates that this highly structured solvent needs to destructure to flow. This is quantified by the entropic contribution that is more important than the enthalpic one in the activation Gibbs energy for viscous flow. In a second part, we demonstrate that maghemite nanoparticles can be dispersed in this complex solvent thanks to a transfer from water. They are well dispersed, interacting through weak repulsions sufficient to stabilize the dispersions in zero magnetic field and under field up to 1350 Oe. This threshold could be increased using smaller particles and reduced polydispersities. These results are similar to those previously obtained in an ionic liquid, EAN and confirm that the mixture ChEG studied here behaves like an ionic liquid. Indeed, a behaviour of concentrated electrolyte solution would not allow the dispersion of NPs which would flocculate due to the screening of electrostatic interactions. The local viscosity experienced by the NPs in ChEG dispersions as well as its temperature variation indicates that the particles locally modify the solvent around the NPs. This could be a local modification of organization or of composition. The results can probably be extended to other oxide nanoparticles, with the same kind of surface groups (hydroxides).

To go further, quantitative macroscopic viscosity measurements, difficult as samples are black and hygroscopic, would be interesting to compare to microscopic viscosities. Also parallel SAXS measurements as a function of temperature would enable to get a clear picture of the local behavior of the samples.

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Supporting Information

* Table : Viscosity and density values for each temperature T and water content x .

* figure SI-1 : Vmol vs water content x , for each temperature T

* figure SI-2 : SAXS scattered intensity $I(Q)$

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