

Intermediate Range Order in Metal–Ammonia Solutions: Pure and Na-Doped Ca–NH₃

Thomas C. Nicholas, Thomas F. Headen, Jonathan. C. Wasse, Christopher. A. Howard, Neal. T. Skipper, and Andrew G. Seel*



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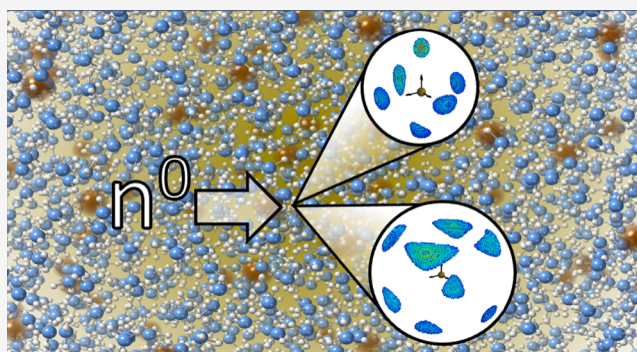
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ABSTRACT: The local and intermediate range ordering in Ca–NH₃ solutions in their metallic phase is determined through H/D isotopically differenced neutron diffraction in combination with empirical potential structure refinements. For both low and high relative Ca concentrations, the Ca ions are found to be octahedrally coordinated by the NH₃ solvent, and these hexammine units are spatially correlated out to lengthscales of ~7.4–10.3 Å depending on the concentration, leading to pronounced ordering in the bulk liquid. We further demonstrate that this liquid order can be progressively disrupted by the substitution of Ca for Na, whereby a distortion of the average ion primary solvation occurs and the intermediate range ion–ion correlations are disrupted.



INTRODUCTION

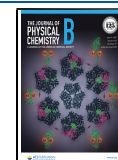
The dissolution of electropositive group I and group II metals in liquid NH₃ yields a rich series of electronic liquids, whose properties are chemically tunable through both the concentration and nature of the parent metal.^{1,2} Ca, as a divalent metal, shares many of the fascinating properties commonly associated with the group I metals when dissolved in NH₃, forming an electrolytic system of solvated electrons at low concentrations and itinerant, metallic liquids as the concentration of Ca is increased.^{1,3} As with other metal–ammonia systems, these different electronic states are not miscible and are separated by a forbidden region around a metal–insulator transition whereby a bulk liquid–liquid phase separation occurs, commonly referred to as the immiscibility region in the literature. Unusually, this immiscibility region is extremely pronounced in the Ca–NH₃ phase diagram, with the electrolytic state only surviving to a maximum Ca concentration of 1.68 mole percent metal (MPM) at the consolute point or a critical temperature of 290 K with the onset of phase separation at <0.1 MPM at 200 K.^{1,4} For comparison, the Na–NH₃ consolute point is 4.12 MPM and 231 K with phase separation beginning at 1 MPM at 220 K, and in Li–NH₃, the consolute point is 4.35 MPM at 210 K with phase separation beginning at 2 MPM at 220 K.⁵ Such early onset and pronounced liquid–liquid phase separation clearly indicate the stability of, or energetic drive toward, the metallic liquid in Ca–NH₃ solutions.

Previous studies have sought to understand the link between the different electronic states of metal–ammonia systems and their liquid structure, whereby the charge of the metal cation, the extent of screening by the itinerant electron, and the secondary solvation effects from the NH₃ solvent can all be varied. Some metal–ammonia systems, including Li–NH₃ and Ca–NH₃, have been shown to possess strong intermediate range ordering between ~6 and 10.5 Å in their metallic phases, whereas other systems such as Na–NH₃ and K–NH₃ do not.^{6–9} Interestingly, only those systems which exhibit such strong ordering in their liquid state are known to crystallize as solid phases at their concentration limit. All other metal–ammonia or metal–amine systems precipitate out the metal upon cooling and do not form solid metal–amine phases. Taking only the examples of the Ca–NH₃ and Na–NH₃ systems, it is notable that, despite their similar ionic radii,¹⁰ Ca–NH₃ possesses distinct intermediate range ordering in the liquid and crystallizes as the expanded metal Ca(NH₃)₆,^{9,11} whereas no such liquid order nor any crystalline phases have been witnessed in Na–NH₃.⁸

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Herein, we use neutron scattering with combined H/D isotopic substitutions and empirical-potential structure refinements¹² to provide a more detailed study of the Ca–NH₃ system in the metallic phase than has been performed previously. Both the local and intermediate solvation environments of the Ca ions are determined alongside Ca–Ca correlations in the liquid phase. We also demonstrate that the intermediate range order of the Ca–NH₃ solution can be successively disrupted by the introduction of Na.

EXPERIMENTAL METHODS

Stoichiometric amounts of mechanically cleaned metal were loaded under argon into a sealed flat-plate null-coherent scattering Ti/Zr container of size 15 mm × 30 mm × 1 mm. The container and sample were attached to a stainless steel gas rig and evacuated to <10^{−5} mbar. Appropriate ratios of isotopically unique ammonia (NH₃/ND₃) were premixed to the exact volume required and cryogenically pumped onto the metal samples. The samples were isolated, warmed, and monitored for homogeneity. No notable sample decomposition occurred as monitored by lack of H₂ evolution.

H/D substitution experiments on the Ca–NH₃ and mixed metal NaCa–NH₃ solutions were performed using the SANDALS and NIMROD diffractometers, respectively, at the ISIS Spallation Neutron and Muon Source, UK.^{13,14}

The sample data were corrected for sample containment, multiple scattering, and incoherent scattering according to standard procedures using the Gudrun package.^{15,16} The resultant total structure factor can be expressed as

$$F(Q) = \sum_{\alpha=1}^n \sum_{\beta=1}^n c_{\alpha} c_{\beta} b_{\alpha} b_{\beta} [S_{\alpha\beta}(Q) - 1] \quad (1)$$

where c_i and b_i are the atomic fraction and bound coherent neutron scattering length of species i , respectively. $S_{\alpha\beta}(Q)$ is the Faber–Ziman partial structure factor, and Q is the magnitude of the scattering vector.

The real space function $G(r)$ corresponding to $F(Q)$ is obtained by replacing $S_{\alpha\beta}(Q)$ with the partial pair distribution functions $g_{\alpha\beta}(r)$ by the signed Fourier transform relationship to give

$$G(r) = \sum_{\alpha=1}^n \sum_{\beta=1}^n c_{\alpha} c_{\beta} b_{\alpha} b_{\beta} [g_{\alpha\beta}(r) - 1] \quad (2)$$

In the current context, $G(r)$ can be written as a sum of three partial pair distribution functions

$$G(r) = c_X^2 b_X^2 [g_{XX}(r) - 1] + 2c_X c_H b_X b_H [g_{XH}(r) - 1] + c_H^2 b_H^2 [g_{HH}(r) - 1] \quad (3)$$

where the composite coherent scattering length b_X and atomic concentration c_X are defined as

$$b_X = \sum_{\alpha \neq H} \frac{c_{\alpha} b_{\alpha}}{c_X}, \quad c_X = \sum_{\alpha \neq H} c_{\alpha} \quad (4)$$

The label H refers to substituted atoms in the sample, and X refers to unsubstituted atomic species. It follows from eq 4 that $c_H = 1 - c_X$. Then, the $g_{HH}(r)$ term in eq 3 is calculated from

$$g_{HH}(r) - 1 = \frac{xG_H(r) + (1-x)G_D(r) - G_{HD}(r)}{c_H^2 (xb_H^2 + (1-x)b_D^2 - b_{HD}^2)} \quad (5)$$

where the subscripts H, D, and HD refer to experiments on the protonated, deuterated, and mixture samples, respectively, and x is the fraction of protonated ammonia in the mixture sample. Then

$$b_{HD} = xb_H + (1-x)b_D \quad (6)$$

The remaining partial pair distribution functions can be calculated in an analogous way to $g_{HH}(r) - 1$ in eq 3

$$g_{XH}(r) - 1 = [G_H(r) - G_D(r) - c_H^2 b_H^2 (g_{HH}(r) - 1) + c_H^2 b_D^2 (g_{HH}(r) - 1)] / [2c_H c_X b_H b_X - 2c_H c_X b_D b_X] \quad (7)$$

$$g_{XX}(r) - 1 = [G_H(r) - 2c_H c_X b_H b_X (g_{XH}(r) - 1) - c_H^2 b_H^2 (g_{HH}(r) - 1)] / c_X^2 b_X^2 \quad (8)$$

Structural models for the liquid systems at 4 and 10 MPM were refined to the respective neutron datasets using the empirical potential structure refinement (EPSR) software.¹² This technique refines a structural model based upon an equilibrated Monte Carlo simulation using seed pair-potentials. These potentials are iteratively modified through the introduction of an empirical potential based upon the difference between simulated and measured, isotopically unique neutron scattering structure factors. Simulation boxes consisted of 6000 molecules within the cubic cell of length 64.588 and 66.674 Å for the 4 and 10 MPM systems, respectively, constructed using the reported densities for the Ca–NH₃ system. The seed Lennard-Jones plus Coulomb potentials used in the refinement are reported in Table 1. The

Table 1. Lennard-Jones Plus Coulomb Parameters for Seed Potentials Used in EPSR Refinements

atom	σ_{aa} (Å)	ϵ_{aa} (kJ mol ^{−1})	q (e)
N	3.420	0.711	−1.020
H	0.0	0.0	0.34− y^a
Ca	2.412	1.380	1.530

^a y is introduced to ensure a net-neutral system, defined as $N_{Ca}q_{Ca}/3N_{NH_3}$ where N_i is the number of species i .

all-atom optimized potential for liquid simulation Lennard-Jones parameters of Jorgensen have been used for NH₃ with an adjustable Coulomb term to ensure net neutrality in the simulations of the itinerant electron systems.¹⁷ The reduced charge of Ca ions was taken as those reported from *ab initio* simulations for single ions in NH₃ solutions¹⁸ and was found to be necessary for subsequent structural refinement to the experimental neutron data.

RESULTS AND DISCUSSION

The experimental neutron total structure factors, $F(Q)$, for metallic Ca–NH₃ solutions at 4 and 20 MPM are presented in Figure 1 for the deuterated, proteated, and mixed proteated–deuterated samples. The sharp, concentration-dependent prepeak at the lowest Q -values is characteristic of those metallic solutions which possess a strong intermediate range order of the solvated Ca²⁺ cationic centers with increasing metal content. Their occurrence at 0.61 and 0.85 Å^{−1} indicate Ca–Ca correlations at ~7.4 and ~10.3 Å in the 10 and 5 MPM solutions, respectively. Our measurements are thus in

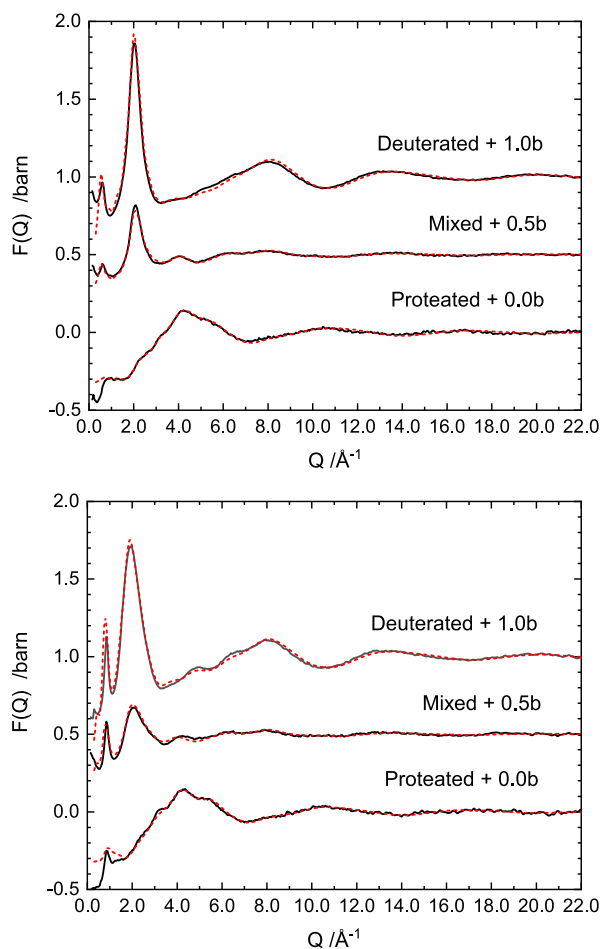


Figure 1. Total structure factor $F(Q)$ data for isotopically unique Ca-NH_3 samples at concentrations 4 MPM (top) and 10 MPM (bottom). Data are offset by a given amount for clarity. From top to bottom: deuterated, mixed, and proteated. EPSR refinements to the data are shown by dashed red lines.

agreement with previous neutron measurements of Ca-NH_3 .^{6,9} It can be seen that the principal diffraction peak in the $2.5\text{--}3.0 \text{ \AA}^{-1}$ region displays considerable broadening with increasing metal concentration, and the feature between 4.0 and 10.0 \AA^{-1} is distinctly structured in the more concentrated Ca-NH_3 solution.

The origin of these features is clearer when looking at the real space transforms for the neutron data. Figures 2 and 3 show the real space correlation functions, $G(r)$, for the 4 and 10 MPM Ca-NH_3 data, respectively, as well as the extracted partial pair distribution functions, $G_{\text{HH}}(r)$, $G_{\text{XH}}(r)$, and $G_{\text{XX}}(r)$ as defined in eqs 5, 7, and 8. It can be seen that the $G_{\text{XH}}(r)$ and $G_{\text{XX}}(r)$ partials are far more structured in the 10 MPM Ca-NH_3 solution than they are at 4 MPM in the $2.5\text{--}6.0 \text{ \AA}$ region, indicating a distinctly ordered solvation environment of the Ca ions. In particular, we can assign the feature at 2.52 \AA in $G_{\text{XX}}(r)$ to the Ca–N distance, being far more pronounced in the 10 MPM solution. Likewise, the narrowing in the most intense feature to 3.2 \AA and the appearance of a peak at 5.0 \AA can be assigned to *cis*- and *trans*-N–N distances, which would be consistent with an octahedral arrangement of NH_3 around Ca. It is interesting to compare these liquid, metallic Ca-NH_3 solutions to solvated, aqueous calcium solutions,¹⁹ whereby the Ca–O distances reported are only slightly shorter at 2.46 \AA

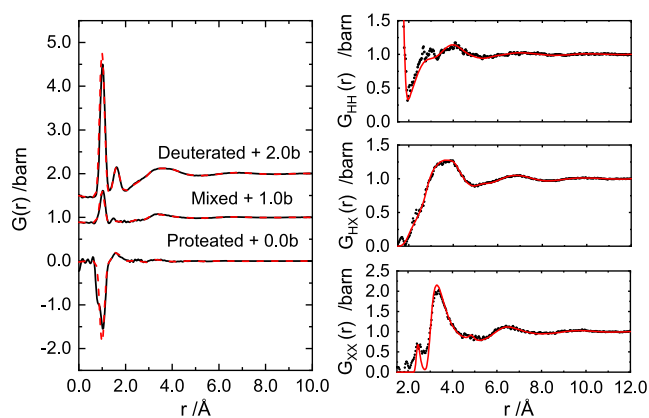


Figure 2. Left: Real space function $G(r)$ for isotopically unique 4 MPM Ca-NH_3 samples. Data are offset by a given amount for clarity. From top to bottom: deuterated, mixed, and proteated. Right: Partial pair distribution function data for 4 MPM Ca-NH_3 . From top to bottom: $G_{\text{HH}}(r)$, $G_{\text{HX}}(r)$, and $G_{\text{XX}}(r)$. EPSR refinements to the data are shown by red lines.

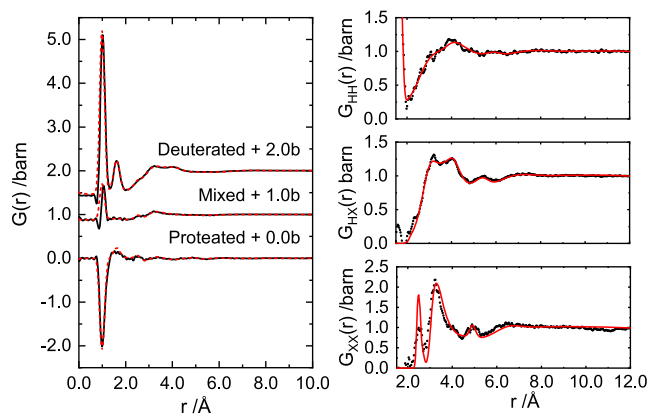


Figure 3. Left: Real space function $G(r)$ for isotopically unique 10 MPM Ca-NH_3 samples. Data are offset by a given amount for clarity. From top to bottom: deuterated, mixed, and proteated. Right: Partial pair distribution function data for 10 MPM Ca-NH_3 . From top to bottom: $G_{\text{HH}}(r)$, $G_{\text{HX}}(r)$, and $G_{\text{XX}}(r)$. EPSR refinements to the data are shown by red lines.

despite the aqueous ions being 8-coordinate as opposed to 6-coordinate for the amines. Turning to the $G_{\text{XH}}(r)$ partials, we see again that the 10 MPM system is more structured with correlations presumably being successive Ca–H, N–H distances.

In order to extract more information from the experimental datasets, individual pair distribution functions were determined from EPSR models for both 4 and 10 MPM systems, with refinements reported alongside the experimental data in Figures 1–3. These structural models, refined as they are to the isotopically unique neutron datasets, are a notable improvement on previous Monte Carlo simulations for the Ca-NH_3 solutions.⁹ The refinements for both 4 and 10 MPM solutions are well described even down to the low- Q region where the relative intensities of the prepeak in $F(Q)$ are matched across datasets, albeit slightly over-fit in the 10 MPM system. Likewise, the full extent of the structuring of the feature between 4.0 and 10.0 \AA^{-1} in the more concentrated Ca-NH_3 solution is not fully captured by the EPSR model. We can see clearly where this discrepancy arises by examining the simulated $G_{\text{XX}}(r)$. The Ca–N distance is correct in the

EPSR refinement, but the feature is sharper than in the measured neutron data, where the intensity is moved into the peak wings. This is most likely due to the insufficiencies of the seed and empirical potentials being too “hard” for a good description of the Ca ions in a metallic solution. Nevertheless, the high level of agreement between our structural models and the neutron data lends credence to our extracted pair distribution functions presented in Figure 4.

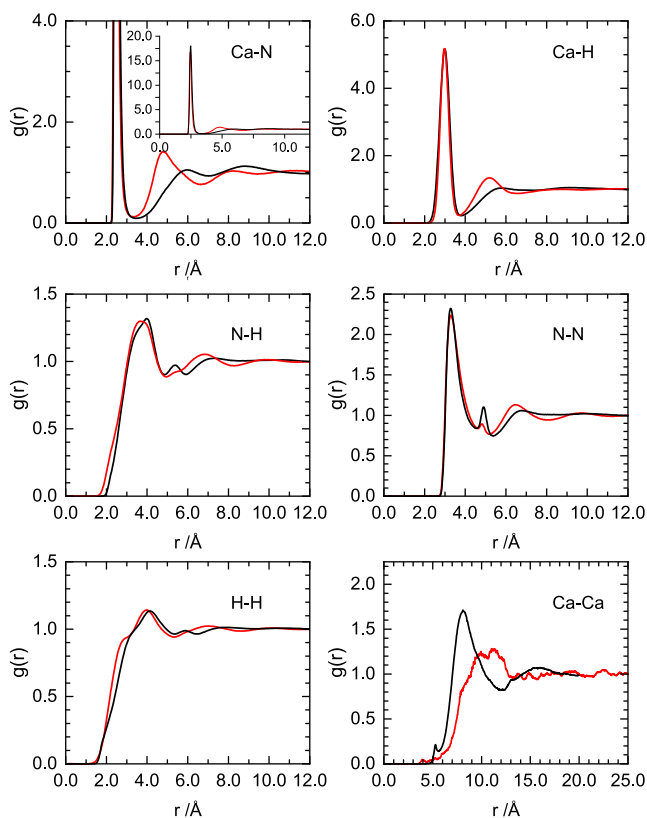


Figure 4. Extracted partial pair radial distribution functions for 4 MPM (red) and 10 MPM (black) Ca–NH₃ solutions. Column 1, from top to bottom: $g_{\text{CaN}}(r)$, $g_{\text{NH}}(r)$, and $g_{\text{HH}}(r)$. Column 2, from top to bottom: $g_{\text{CaN}}(r)$, $g_{\text{NH}}(r)$, and $g_{\text{HH}}(r)$.

The pair distribution functions extracted from the EPSR model allow us to confirm our assignment of the Ca–N and N–N distances from the measured neutron data and the origin of the prepeak as being due to Ca–Ca correlations. The structuring of the measured $G_{\text{XH}}(r)$ partials in the 10 MPM solution can now be interpreted further as the increasing contribution from the Ca–H correlations at 3.0 Å as the Ca concentration is increased. Only a minor low- r shoulder is seen in the N–H partials, in agreement with the large disruption of any hydrogen bonding in these metallic systems, a commonality of all measured metal–amine systems in their metallic phases. The EPSR models also allow us to determine correlations which are only a minor contribution to the experimental data, such as the secondary solvation of Ca(NH₃)₆ in the 4 MPM solution. Clear secondary and tertiary solvation distances are found at 4.8 and 8.2 Å.

In order to build up a three-dimensional picture of the Ca–NH₃ system, Figure 5 details the angular distributions and spatial probability density maps extracted from the refined structural models for Ca–NH₃ solutions. As expected from our assignments of the experimental data, the inner coordination

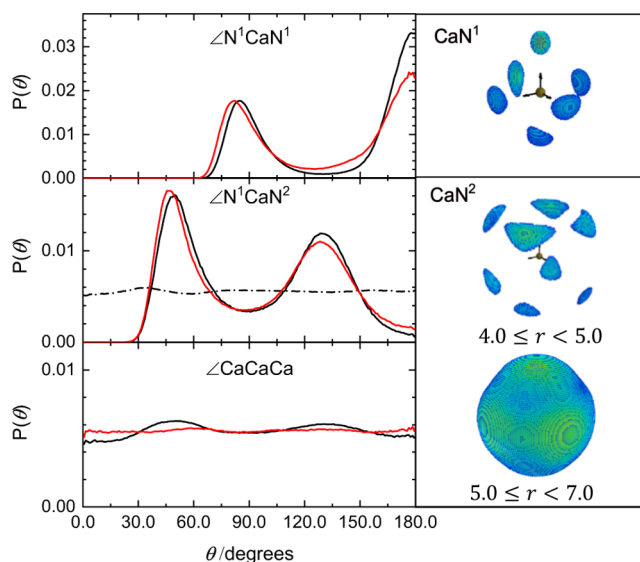


Figure 5. Left: Angular probability densities for 4 MPM (red) and 10 MPM (black) Ca–NH₃ solutions. Labels indicate primary solvation environment around Ca (N¹, 2 ≤ r ≤ 3 Å), secondary solvation (N², 4 ≤ r ≤ 7 Å) where the full and dashed lines for the 10 MPM system are integrated in the ranges 4 ≤ r ≤ 5 and 5 ≤ r ≤ 7 Å, respectively, and Ca ion–ion orientations. Right: Corresponding spatial density maps for N atoms around a central Ca ion in the 10 MPM system.

sphere (corresponding to the sharp first peak in $g_{\text{CaN}}(r)$ in Figure 4) shows an octahedral arrangement of ammonia nitrogen atoms about the central Ca. This solvation is more regular for the 10 MPM system, with the 4 MPM system having a slightly less well-defined octahedral geometry as demonstrated with the broadening of angular distributions for NH₃ coordinated trans to each other. The second coordination shell (corresponding to the broader $g_{\text{CaN}}(r)$ peak in the range of 4.0–7.0 Å) contains those NH₃ molecules not directly bound to the Ca ions in solution and demonstrate a drastic change between those located up to ~5 Å and those beyond. The second peak in $g_{\text{CaN}}(r)$ for the 4 MPM solutions occurs at 4.8 Å, where there is only a minor contribution in the 10 MPM system. This can be assigned to the hydrogen-bonded, secondary solvation of the Ca centers and can be seen in Figure 5 corresponding to a facial approach to the primary Ca(NH₃)₆ species. The 10 MPM system exhibits its secondary solvation maximum in $g_{\text{CaN}}(r)$ at 5.9 Å, at which distance there is very little angular dependence in the spatial density. This demonstrates that neighboring Ca(NH₃)₆ units are only weakly correlated in terms of their relative orientations despite the strong radial correlation between Ca ions in the metallic solutions. While in the 4 MPM solutions, there is no angular correlation in the distribution of Ca ions, in the 10 MPM solution, we do find a slight dominance for a facial approach of one Ca(NH₃)₆ unit to the next, consistent with the *bcc* arrangement of the solid calcium hexammine structure, Ca(NH₃)₆, formed at the pseudo-eutectic point whereby cooling any metallic Ca–NH₃ solution to 185 K excess NH₃ crystallizes out leaving only the expanded metal.^{20,21} This expanded-metal hexammine crystallizes in the *Im3m* space group with a lattice constant, $a_0 = 9.12$ Å, and an octahedral orientation of NH₃ molecules around each Ca atom.^{11,20} Bond distances in the crystal have been determined as $r(\text{Ca–N}) = 2.56$ Å and $r(\text{Ca–D}) = 3.12$ Å, correlations which we have now

shown to persist into the liquid state, even upon further dilution by additional NH_3 as in the 4 MPM solution.

We have previously suggested that for a metal–amine system to crystallize into an expanded metal upon cooling, there must be a substantial order already present in the liquid both in terms of primary solvation and intermediate range order of the metal centers. This leads to the stability of the pseudo-eutectic and prevents the formation of crystalline NH_3 and metal. Our findings for the Ca-NH_3 system further strengthen this viewpoint and certainly provide a solution-phase limit in the future search for new, crystalline expanded metals.

We can investigate the manner in which the structural order in the liquid Ca-NH_3 system can be disrupted by examining the effect of progressive substitution of Ca by Na. The ionic radii of Ca^{2+} and Na^+ are similar, and both have been determined to be octahedrally coordinated in aqueous salt solutions, but in the case of Na^+ , there is some uncertainty in this, with the experimentally determined coordination numbers ranging from 4 to 8.^{22,23} We should also note that when considering NaCa-NH_3 solutions in their metallic phase, despite maintaining the concentration of metal ions (MPM), the density of conduction electrons in the solution (sometimes referred to as mole-percent electrons) can vary up to a factor of 2 due to the difference in valence between Na and Ca. This conduction electron density is believed to lie predominantly between solvated ion units in metal–amine solutions, and we can thus determine the effect on the liquid structure of varying this electron density through control of the Na/Ca content.

Figure 6 reports the experimental neutron structure factors for the 10 MPM, deuterated NaCa-NH_3 system at various

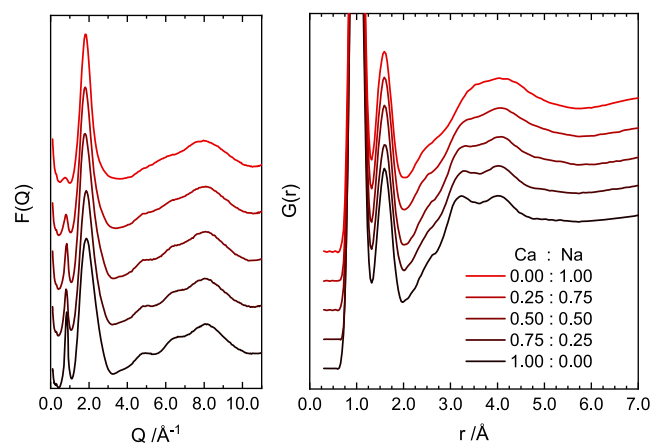


Figure 6. Left: Neutron total structure factor $F(q)$ data for Na-doped 10 MPM Ca-NH_3 solutions. Data are offset for clarity. Right: Corresponding $G(r)$ data.

Na/Ca ratios. We can clearly see that the pronounced prepeak in Ca-NH_3 rapidly decreases in intensity as the Na content increases, indicating that ion–ion correlations are decreasing, and both the width of the principal peak and structuring of the feature from 4.0 to 10.0 Å^{-1} decrease. This would indicate a disruption of the octahedral arrangement of the solvated metal ions, which can be confirmed by examining the equivalent data in real space. Interestingly, while the M-NH_3 correlations are still found at ~ 2.5 Å , we see a slight shift to shorter bond lengths as the Na content increases, despite the lower charge of these ions, and there is a progressive disruption of the structure

in the 3.0–5.0 Å range. This shorter bond is consistent with the Na–N distances previously reported for pure Na-NH_3 solutions in their metallic phase.⁸ It is likely that the longer Ca–N distances in Ca-NH_3 is a reflection of the highly structured $\text{Ca}(\text{NH}_3)_6$ units being more “molecular” in nature than the Na-NH_3 analogue. It is clear that the incorporation of Na ions into the Ca-NH_3 solution disrupts the local solvation structure of the metal ions, with the distinctly octahedral coordination of Ca being lost upon substitution by Na. This in turn leads to a loss of intermediate range order in the mixed-ion solutions.

CONCLUSIONS

Despite its molecular nature and the large concentration range over which a metallic phase occurs, the liquid Ca-NH_3 system is structurally quite simple. There is a strong local ordering whereby the octahedral coordination of Ca by NH_3 comprise the solvated ion cores for both the most dilute and more concentrated itinerant electron liquids. The approach of additional NH_3 is toward the octahedral faces, being driven by the formation of hydrogen bonding interactions with the solvated ion cores. Beyond this, further NH_3 molecules still exist in distinct solvation shells but are spatially isotropic. It is striking to note that even at low metal concentrations of 4 MPM, a significant intermediate range order exists in these solutions, with the solvated ion cores being correlated out to >10 Å . This becomes more pronounced and with a shift to a shortened distance of 7.4 Å for the 10 MPM solution. As we have previously noted, this structuring in the liquid state seems to be a feature of metal–amine systems, which undergo crystallization at their concentration limit, and is significantly reduced or absent in those which do not.²⁴

Furthermore, we have showed that the miscibility of Na– and Ca– NH_3 enables the local and intermediate structure of the bulk liquid to be controlled as a function of the conduction electron density. Sequential disruption of the primary solvation environment upon incorporation of Na is evident from the neutron scattering data, which in turn is reflected in the intermediate range, ion–ion correlations being reduced. Whether this has an effect on the electronic properties of the liquids, or indeed one which can be carried across into any crystalline material, is yet to be determined. It is now clear that, at the very least, the liquid structure for metal– NH_3 and other metal–amine systems is amenable to chemical control.

AUTHOR INFORMATION

Corresponding Author

Andrew G. Seel – Department of Physics and Astronomy,
University College London, London WC1E 6BT, U.K.;
orcid.org/0000-0003-0103-8388; Email: a.seel@ucl.ac.uk

Authors

Thomas C. Nicholas – Department of Chemistry, Inorganic Chemistry Laboratory, University of Oxford, Oxford OX1 3QR, U.K.

Thomas F. Headen – ISIS Spallation Neutron and Muon Source, STFC Rutherford Appleton Laboratory, Didcot, Oxfordshire OX11 0QX, U.K.; orcid.org/0000-0003-0095-5731

Jonathan. C. Wasse – Department of Physics and Astronomy, University College London, London WC1E 6BT, U.K.

Christopher. A. Howard – Department of Physics and Astronomy, University College London, London WC1E 6BT, U.K.; orcid.org/0000-0003-2550-0012

Neal. T. Skipper – Department of Physics and Astronomy, University College London, London WC1E 6BT, U.K.; orcid.org/0000-0003-2940-3084

Complete contact information is available at:
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Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) Thompson, J. C. *Electrons in Liquid Ammonia*; Oxford University Press: Oxford, 1976.
- (2) Zurek, E.; Edwards, P. P.; Hoffmann, R. A Molecular Perspective on Lithium–Ammonia Solutions. *Angew. Chem., Int. Ed.* **2009**, *48*, 8198–8232.
- (3) Catterall, R. Metal–Ammonia Solutions. *Adv. Phys.* **1969**, *18*, 665.
- (4) Thompson, J. C.; Teoh, H.; Antoniewicz, P. R. Coexistence of Liquid Phases in Calcium–Ammonia Solutions. *J. Phys. Chem.* **1971**, *75*, 399–405.
- (5) Lodge, M. T. J. H.; Cullen, P.; Rees, N. H.; Spencer, N.; Maeda, K.; Harmer, J. R.; Jones, M. O.; Edwards, P. P. Multielement NMR Studies of the Liquid-Liquid Phase Separation and the Metal-to-Nonmetal Transition in Fluid Lithium- and Sodium–Ammonia Solutions. *J. Phys. Chem. B* **2013**, *117*, 13322–13334.
- (6) Chieux, P.; Bertagnolli, H. Deuterated Liquid Ammonia and ^7Li - 4NH_3 Solution. A Neutron Scattering Investigation. *J. Phys. Chem.* **1984**, *88*, 3726–3730.
- (7) Wasse, J. C.; Hayama, S.; Skipper, N. T.; Benmore, C. J.; Soper, A. K. The structure of saturated lithium- and potassium-ammonia solutions as studied by using neutron diffraction. *J. Chem. Phys.* **2000**, *112*, 7147–7151.
- (8) Wasse, J. C.; Hayama, S.; Masmanidis, S.; Stebbings, S. L.; Skipper, N. T. The structure of lithium-ammonia and sodium-ammonia solutions by neutron diffraction. *J. Chem. Phys.* **2003**, *118*, 7486–7494.
- (9) Wasse, J. C.; Howard, C. A.; Thompson, H.; Skipper, N. T.; Delaplane, R. G.; Wannberg, A. The structure of calcium-ammonia solutions by neutron diffraction. *J. Chem. Phys.* **2004**, *121*, 996.
- (10) Lodge, M. T. J. H. Multielement NMR Studies of the Liquid-Liquid Phase Separation and the Metal-to-Nonmetal Transition in Fluid Lithium- and Sodium–Ammonia Solutions. *J. Phys. Chem. B* **2013**, *117*, 13322–13334.
- (11) Damay, P.; Leclercq, F.; Chieux, P. Geometry of the ND_3 group in a metallic $\text{Ca}(\text{ND}_3)_6$ compound and in solid and liquid deuterioammonia as measured by neutron scattering. *Phys. Rev. B: Condens. Matter Mater. Phys.* **1990**, *41*, 9676–9682.
- (12) Soper, A. K. Partial structure factors from disordered materials diffraction data: An approach using empirical potential structure refinement. *Phys. Rev. B: Condens. Matter Mater. Phys.* **2005**, *72*, 104204.
- (13) <http://www.isis.stfc.ac.uk/instruments/sandals>.
- (14) <http://www.isis.stfc.ac.uk/instruments/nimrod>.
- (15) Soper, A. K. GudrunN and GudrunX: programs for correcting raw neutron and X-ray diffraction data to differential scattering cross section. *RAL Reports*, 2011.
- (16) Soper, A. K. The Radial Distribution Functions of Water as Derived from Radiation Total Scattering Experiments: Is There Anything We Can Say for Sure? *ISRN Phys. Chem.* **2013**, *2013*, 1–67.
- (17) Rizzo, R. C.; Jorgensen, W. L. OPLS All-Atom Model for Amines: Resolution of the Amine Hydration Problem. *J. Am. Chem. Soc.* **1999**, *121*, 4827–4836.
- (18) Chaban, V. V.; Prezhdo, O. V. Electron Solvation in Liquid Ammonia: Lithium, Sodium, Magnesium, and Calcium as Electron Sources. *J. Phys. Chem. B* **2016**, *120*, 2500–2506.
- (19) Bruni, F.; Imberti, S.; Mancinelli, R.; Ricci, M. A. Aqueous Solutions of Divalent Chlorides: Ions Hydration Shell and Water Structure. *J. Chem. Phys.* **2012**, *136*, 064520.
- (20) Mammano, N.; Sienko, M. J. An X-ray study of the alkaline earth hexaamines at 77 K. *J. Solid State Chem.* **1970**, *1*, 534–535.
- (21) Bridges, R.; Ingle, A. J.; Bowen, D. E. Sound Velocities in Concentrated Metal–Ammonia Solutions. *J. Chem. Phys.* **1970**, *52*, 5106–5111.
- (22) Ohtaki, H.; Radnai, T. Structure and Dynamics of Hydrated Ions. *Chem. Rev.* **1993**, *93*, 1157–1204.
- (23) Mancinelli, R.; Botti, A.; Bruni, F.; Ricci, M. A.; Soper, A. K. Hydration of Sodium, Potassium, and Chloride Ions in Solution and the Concept of Structure Maker/Breaker. *J. Phys. Chem. B* **2007**, *111*, 13570–13577.
- (24) Seel, A. G.; Holzmann, N.; Imberti, S.; Bernasconi, L.; Edwards, P. P.; Cullen, P. L.; Howard, C. A.; Skipper, N. T. Solvation of Na^- in the Sodide Solution, $\text{LiNa}-10\text{MeNH}_2$. *J. Phys. Chem. B* **2019**, *123*, 5337–5342.