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Unravelling the complex speciation of halozincate ionic liquids using X-ray spectroscopies and calculations

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Abstract

Using a combination of liquid-phase experimental X-ray spectroscopy experiments and small-scale calculations we have gained new insights into the speciation of halozincate anions in ionic liquids (ILs). Both core and valence X-ray photoelectron spectroscopy (XPS) were performed directly on the liquid-phase ILs, supplemented by Zn 1s X-ray absorption near edge structure (XANES) spectroscopy. Density functional theory (DFT) calculations were carried out on both 1- and 2- halozincate anions, in both a generalised solvation model SMD (Solvation Model based on Density) and the gas phase, to give XP spectra and total energy differences; time-dependent DFT was used to calculate XA spectra. Speciation judgements were made using a combination of the shape and width of experimental spectra, and visual matches to calculated spectra. For 2- halozincate anions, excellent matches were found between experimental and calculated XP spectra, clearly showing that only 2- halozincate anions were present at all zinc halide mole fraction, x , studied. At specific x (0.33, 0.50, 0.60) only one halozincate anion was present; equilibria of different halozincate anions at those x were not observed. All findings show that chlorozincate anion and bromozincate anion speciation matched at the same x . Based on the results, predictions are made of the halozincate anion speciation for all x up to 0.67. Caution is advised when using differences in calculated total energies obtained from DFT to judge halozincate anion speciation, even when the SMD was employed, as predictions based on total energy differences did not always match the findings from experimental and calculated spectra. Our findings clearly establish that the combination of high-quality experimental data from multiple spectroscopies and a wide range of calculated structures are essential to have high confidence in halozincate anion speciation identification.

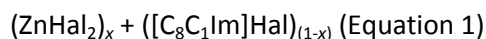
1. Introduction

Halozincate ionic liquids (ILs), formed by ZnHal_2 (Hal = halide) added to a halide anion-containing IL (Equation 1, e.g. 1-octyl-3-methylimidazolium chloride, $[\text{C}_8\text{C}_1\text{Im}]\text{Cl}$),¹ have potential applications as catalysts,² in zinc electrodeposition,³⁻⁵ in zinc batteries⁶⁻⁸ extraction/separation⁹⁻¹³ and preparing antibacterial surfaces^{14, 15}. The variation in speciation (*i.e.* zinc anions present in solution) with increasing mole fraction of ZnHal_2 , x , in halozincate ILs potentially allows fine control of both properties (*e.g.* mass transport) and reactivity (*e.g.* catalysis, electrodeposition). Lewis acidity and thermal stability increase with increasing x ,^{16, 17} and anion-cation interaction strength and cost decrease with increasing x .¹⁷⁻¹⁹ Both monomeric and oligomeric halometallate anions can be formed for $\text{ZnHal}_2 + [\text{C}_8\text{C}_1\text{Im}]\text{Hal}$, with oligomeric halometallate anions composed of multinuclear metals and bridging halides; oligomeric halometallate anions are not formed for all halometallate ILs, *e.g.* $\text{FeCl}_2 + \text{Cl}^-$ which forms solid precipitate at $x > 0.33$.²⁰ The primary halide ligand studied has been chloride,¹ with some focus on bromide. The combination of appealing properties and the ability to tune those



properties by varying the composition through sensible choices of both x and halide identity makes halozincate particularly attractive.

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Understanding halozincate speciation is crucial, as it allows informed selection of x and halide. The option to test all x and halide combinations to find the optimum composition for a particular application is not feasible, given the large range of possible x and more than one possible halide. Furthermore, calculated dissociation energies suggest that terminal and bridging halides in halozincate anions have different Zn-Hal bond strengths,¹⁷ demonstrating that knowledge of speciation is crucial for interpretation of reactivity data in particular. For example, knowledge of speciation would enable determination of both catalytic and electrodeposition mechanisms, allowing the design of better catalysts and optimisation of reactivity.

Zn is spectroscopically quiet,²¹ with a dearth of techniques available that probe liquid phase speciation. Spectroscopies that are suitable for open shell halometallate complexes are unsuitable for probing halozincate anion speciation, *e.g.* electron paramagnetic resonance (EPR) and UV-VIS spectroscopy. Indirect methods have been used to study zinc speciation, *e.g.* bulk property measurements. Mass spectrometry has been widely used, although questions have been raised about the suitability of using both gas phase and solid phase techniques to probe liquid phase speciation,^{1, 22} especially since most 2- halometallate anions are unstable in the gas phase due to strong electrostatic repulsion of the excess electrons.²³ Raman spectroscopy, core X-ray photoelectron spectroscopy (XPS), X-ray absorption near edge structure (XANES) spectroscopy and extended X-ray absorption fine structure (EXAFS) have been recommended as the best techniques for probing halozincate speciation.¹ Furthermore, calculated anion total energy differences from small-scale density functional theory (DFT) calculations have been used to judge speciation for halobismuthate anions (Gibbs free energies were used)²⁴ and halozincate anions (sum of electronic and zero-point energies were used, although the values were given as dissociation energies)¹⁷. A full comparison of spectroscopic and calculated structures has not yet been made for halozincate anions, especially consideration of both 1- and 2- halozincate anions (see Figure 1 for 2- halozincate anions and ESI Figure S1 for 1- halozincate anions).

For Zn^{2+} cations, there is no ligand field stabilisation energy, so the coordination number is primarily determined by the size of the coordinating ligands.²⁵ Zinc is usually tetrahedrally coordinated (although in aqueous solution Zn^{2+} is coordinated to six water molecules),²⁶ which is expected to be driven by the four vacant orbitals of the Zn^{2+} cation ($4s + 4p^3$).²⁷ Therefore, one would expect the same coordination number for all chlorozincate anions and the same coordination number for all bromozincate anions, *i.e.* for all x . A strong case has been made based on Raman spectroscopy and core XPS that halozincate anions are always 2-.^{16, 18} However, a case was made using gas phase mass spectrometry that 1- chlorozincate anions were always formed.²⁸⁻³⁰ Two pertinent observations. Firstly, no one has claimed to have found both 1- and 2- chlorozincate anions for the same sample, indicating essentially two separate groups of findings. Secondly, when calculations were performed on 1- anions it would appear that 2- halozincate anions were not considered, so these calculations should not be used to judge speciation. Therefore, the speciation of halozincate ILs at different x remains an open question.

To quote the excellent review by Swadźba-Kwaśny and co-workers: "The defining characteristic of chlorometallates is the presence of complex anionic equilibria".¹ For halometallate ILs it is possible to form just one halometallate anion at certain x (*e.g.* for halozincates at $x = 0.50$ it is possible to form just $[\text{Zn}_2\text{Cl}_6]^{2-}$) but equilibria of more than one halozincate anion may also form at these x (*e.g.* at $x = 0.50$ $[\text{Zn}_2\text{Cl}_6]^{2-}$ and $[\text{ZnCl}_3]^-$ or $[\text{ZnCl}_4]^{2-}$ and $[\text{Zn}_3\text{Cl}_8]^{2-}$ could potentially form). For $\text{AlCl}_3 + \text{Cl}^-$ multiple



chloroaluminate anions have been found at one x , *e.g.* at $x = 0.67$ both $[\text{Al}_2\text{Cl}_7]^-$ and $[\text{Al}_3\text{Cl}_{10}]^-$ (Figure 8b),^{1, 31} demonstrating that an equilibrium occurs; an excellent summary for $\text{AlCl}_3 + \text{Cl}^-$ is given in reference ¹. View Article Online
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The speciation for $x = 0.50$ is still not settled; does $2 \times \text{ZnCl}_2 + 2 \times \text{Cl}^-$ form $[\text{Zn}_2\text{Cl}_6]^{2-}$ or $2 \times [\text{ZnCl}_3]^-$?²² Using extended X-ray absorption fine structure (EXAFS) data at $x = 0.50$, the coordination number of Zn was 2.9,³² suggesting the formation of $[\text{ZnCl}_3]^-$. Both fast atom bombardment and electrospray ionisation mass spectrometry studies came to the same conclusion of $[\text{ZnCl}_3]^-$ speciation.^{28, 30} Conversely, Raman spectroscopy and core XPS concluded that $[\text{Zn}_2\text{Cl}_6]^{2-}$ was the species at $x = 0.50$.^{16, 18} Calculations have been performed on both $[\text{ZnCl}_3]^-$ ^{33, 34} and $[\text{Zn}_2\text{Cl}_6]^{2-}$.¹⁷

The speciation for $x = 0.67$ is also still not settled; does $4 \times \text{ZnCl}_2 + 2 \times \text{Cl}^-$ form $[\text{Zn}_4\text{Cl}_{10}]^{2-}$ or $2 \times [\text{Zn}_2\text{Cl}_5]^-$?²² Using extended X-ray absorption fine structure (EXAFS) data at $x = 0.67$, the coordination number of Zn was found to be 2.6,³² suggesting the formation of $[\text{Zn}_2\text{Cl}_5]^-$ with two terminal Cl and one bridging Cl for both Zn centres. Using fast atom bombardment mass spectrometry at $x = 0.67$, a range of 1- anions were observed: $[\text{ZnCl}_3]^-$, $[\text{Zn}_2\text{Cl}_5]^-$, $[\text{Zn}_3\text{Cl}_7]^-$ and high mass clusters.^{28, 29} Electrospray ionisation mass spectrometry experiments reported for $x > 0.50$ showed $[\text{Zn}_3\text{Cl}_7]^-$ was present.³⁰ Furthermore, $[\text{Zn}_2\text{Cl}_5]^-$ was assumed to be the anion in ammonia absorption experiments underpinning their use in absorption refrigerators and heat pumps.¹³ Conversely, Raman spectroscopy concluded that $[\text{Zn}_4\text{Cl}_{10}]^{2-}$ was the species at $x = 0.67$.¹⁶ In the solid state, a supertetrahedron, $[\text{Zn}_4\text{Cl}_{10}]^{2-}$, has been found.³⁵ Calculations have been performed on both $[\text{Zn}_2\text{Cl}_5]^-$ (with three bridging Cl and one terminal Cl on both Zn centres)³⁶ and $[\text{Zn}_4\text{Cl}_{10}]^{2-}$.¹⁷

Given that both $[\text{ZnCl}_3]^-$ ^{30, 32} and $[\text{ZnCl}_4]^{2-}$ ^{16, 18, 37} are apparently stable, the speciation for $x = 0.33$ is also not settled; does $1 \times \text{ZnCl}_2 + 2 \times \text{Cl}^-$ form $[\text{ZnCl}_4]^{2-}$ or $1 \times [\text{ZnCl}_3]^- + 1 \times \text{Cl}^-$?²² An interesting property of chlorozincates is that the Lewis acidity (measured using the Gutmann-Beckett method) has a step-change at $x = 0.33$, which indicates a change in speciation at $x = 0.33$.¹⁶

It is still unclear whether multiple halozincate anions exist in equilibrium at one x where it is possible that just one anion forms. For halozincates, mass spectrometry at $x = 0.67$ suggested an equilibrium of $[\text{ZnCl}_3]^-$, $[\text{Zn}_2\text{Cl}_5]^-$, $[\text{Zn}_3\text{Cl}_7]^-$ and high mass clusters,²⁹ but this finding is doubtful given the concerns over the reliability of mass spectrometry for halozincate speciation²².

Element-specific electronic structure can be probed using core XPS and XANES; a major advantage for studying halozincate speciation is the ability to study more than one element, *i.e.* both zinc and the halide can be studied independently. Core state-specific binding energies, E_B , from XPS can capture the presence of different electronic environments,³⁸⁻⁴⁰ including for chlorozincates¹⁸. XANES can also be used to determine structure using core state to unoccupied valence state absorption energies.⁴¹ The ratio of peak areas from core XPS gives the relative abundance of each of those species, as there is no selection rule for core XPS at the same edge for the same element, *e.g.* two Cl $2p_{3/2}$ peaks in a 3:2 ratio means a 3:2 ratio of those two Cl-containing species. XPS can be a surface sensitive technique, but for ILs studied using laboratory XPS (*i.e.* at $h\nu = 1486.6$ eV) and under a normal emission angle, signals reflect bulk IL composition, *i.e.* no surface sensitivity has been observed.⁴² Spin-orbit coupling for any orbital that is not an s orbital (*e.g.* Zn 2p, Cl 2p, Br 3d) contributes two components for each electronic environment in a sample in a predictable peak area ratio and peak splitting, *e.g.* for Cl 2p the peak area ratio is 2:1 with $\Delta E_B(\text{Cl } 2p_{3/2} - \text{Cl } 2p_{1/2}) = 1.60$ eV. The timescales of XPS photoemission and XANES are both fs,⁴³ meaning nuclear motion during the core photoemission/photoexcitation process will not contribute to energy differences; if more than one halozincate species is in equilibrium in the IL, both species will contribute to the spectra and the major challenge for detection is sufficient energy separation of the contributions from different halozincate species. For XPS, the core-state peak full width at half maximum (FWHM) is dependent on the core-state studied (each core-state has a



fixed width contribution due to lifetime broadening⁴⁴), instrumental factors (which are very similar for modern XPS apparatus), vibrational contributions, and the sample speciation.⁴⁵⁻⁴⁸ Molecules and ions in liquids give significantly larger measured FWHM than metal single crystals such as Pt due to their greater disorder.⁴⁹ On a microscopic level, there are a range of different solvation environments for Cl⁻ solvated by [C₈C₁Im]⁺ in the liquid phase, and these different solvation environments contribute to the XPS FWHM;¹⁹ on a macroscopic level, [C₈C₁Im]Cl and [C₈C₁Im]Br both present one halide electronic environment in the liquid phase. Bridging and terminal chlorine atoms have been observed using XPS for both halometallate solids⁵⁰ and chlorozincate anions in ILs.¹⁸ Valence XPS has not been used to probe halometallate speciation for ILs to date.²⁴

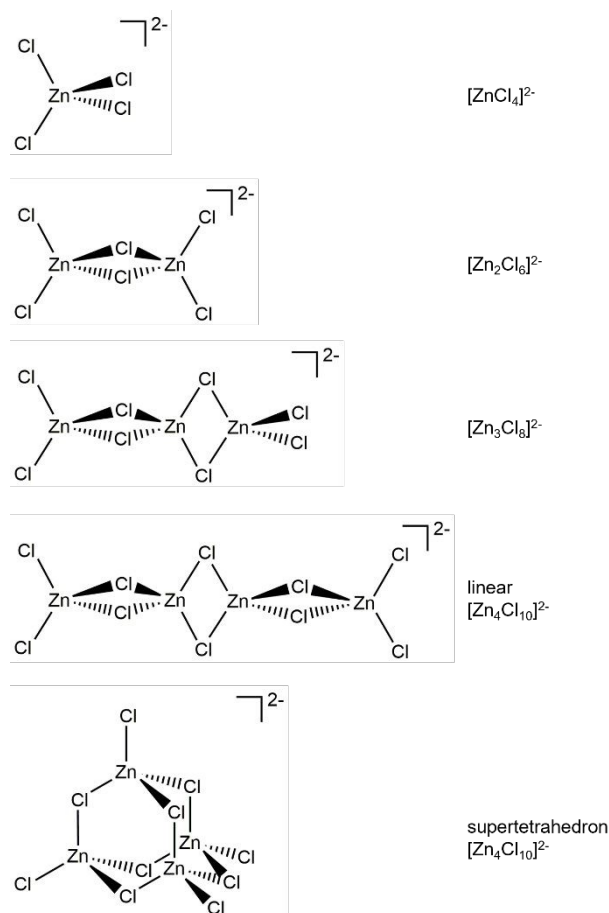


Figure 1. 2- chlorozincate structures. Bromozincate anions would be the same apart from Br replacing Cl. See ESI Figure S1 for 1- chlorozincate structures.

In this article, we report a comprehensive study of the speciation of halozincate anions in ILs. We intend to primarily answer four questions. (i) What is the speciation at specific x where only one halozincate anion may be present? (ii) What contribution does an equilibria of different halozincate anions play in speciation? (iii) Can small-scale DFT calculations capture halozincate speciation? (iv) What is the effect of chloro versus bromo ligands for halozincate speciation? We answered these questions using a combination of core XPS, valence XPS, Zn 1s high energy resolution fluorescence detected (HERFD)-XANES spectroscopy, DFT calculations for XPS (both core and valence) and total energies, time-dependent DFT calculations for Zn 1s XANES spectroscopy. We calculated both 2- and 1- halozincate anions.



2. Methods

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2.1. Experimental

XP spectra for 20 ILs are used in this paper. Core and valence XP spectra were published for 16 ILs in the ESI of reference⁵¹ and for three ILs in the ESI of reference¹⁹. XP spectra are published for the first time for $x = 0.43$ where $(\text{ZnCl}_2)_x + ([\text{C}_8\text{C}_1\text{Im}]\text{Cl})_{(1-x)}$. $E_{\text{B}}(\text{core})$ for all 20 ILs are published for the first time here (ESI Table S4).

The halozincate IL samples (~ 1 drop) were mounted between two pieces of Kapton tape and were kept in place by a plastic O-ring. We used the Si(111) monochromator crystal cut to select the incident energy.⁵² Zn 1s HERFD-XANES measurements were taken using the I20 X-ray emission spectrometer⁵² equipped with three Ge(555) analyser crystals, set at an emission energy of 9572 eV to capture the $K\beta_{1,3}$ ($3p \rightarrow 1s$) X-ray emission.⁵³ The spectrometer was calibrated using a Zn foil, measuring the $K\beta_{1,3}$ line with the incident energy tuned +100 eV from the Zn 1s absorption edge. HERFD-XANES measurements offer better spectral resolution than conventional XANES and are ideally suited for probing unoccupied states.⁵⁴

We emphasise that all XPS and XANES measurements discussed in this article were made at room temperature.

2.2. Calculations

DFT calculations were performed using Gaussian 16 (version C.01).⁵⁵ All zinc halide structures were optimised under no symmetry constraints and confirmed as minima via vibrational analysis employing the long-range corrected ωB97XD functional⁵⁶⁻⁵⁹ in combination with the quadruple zeta def2-QZVPP basis set (see ESI Section 12 for comparisons to data calculated using the triple zeta def2-QZVPP basis set). Several of the initial starting structures were taken from previous work.¹⁷ To account for solvent effects, the SMD (Solvation Model based on Density) was used, specifically the $[\text{C}_4\text{C}_1\text{Im}][\text{PF}_6]$ parameters (relative permittivity, $\epsilon_r = 11.40$; refractive index, $n = 1.4090$; surface tension, $\gamma = 0.266$ cal mol⁻¹ Å²; Abraham basicity, $\beta = 0.216$).⁶⁰ Moreover, optimisation convergence criteria were set to 10^{-11} on the density matrix and 10^{-9} on the energy matrix, and the numerical grid was improved from the default using the keywords (int=SuperFineGrid) which gives a pruned (optimised) grid of 250 radial shells and 974 angular points per shell. Vibrational frequencies and zero-point vibrational energy corrections (ZPE) were attained using the harmonic approximation.

The procedure to convert calculated E_{B} into calculated XP spectra is explained in ESI Section 4. Spin-orbit coupling contributions were added for Cl 2p, Br 3d and Zn 2p, with the parameters given in ESI Table S3.

Single point calculations were computed using ORCA (v 5.0.3).⁶¹ Electronic structure was calculated with time-dependent DFT for Zn 1s XANES, using the $\omega\text{B97X-D3BJ}$ functional,⁶² the ZORA-def2-QZVPP basis set⁵⁹ and SARC/J auxiliary basis set⁵⁸. Scalar relativistic effects were taken into account by applying the zeroth order regular approximation (ZORA).⁶³ In all cases, electric-dipole, magnetic-dipole, and quadrupole contributions were allowed in spectral calculations. Time-dependent DFT calculations were performed with the Tamm-Dancoff approximation (TDA) applied. For the XANES calculation, the Zn 1s orbital was excited into all virtual unoccupied molecular orbitals to mimic the Zn 1s K-edge XANES. Calculated XANES spectra were shifted by -8.9 eV to align with the experimental absorption energies. All XANES spectra were generated by convoluting the computed energies and oscillator strengths with Gaussian functions with a full width at half maximum (FWHM) of 2.5 eV.



2.3. Fitting/Analysis

All experimental XP spectra were fitted using CASAXPS™ software.⁶⁴ Fitting was carried out using a Shirley background and GL30 line shapes (70% Gaussian, 30% Lorentzian). Peak constraints used are outlined in the ESI Section 2. Relative sensitivity factors from reference⁶⁵ were used to ensure the experimentally-derived stoichiometries matched the nominal stoichiometries.

All XPS $E_B(\text{core,exp.})$ were shifted relative to $E_B(\text{C}_{\text{alkyl}}\ 1s,\text{exp.}) = 285.00\ \text{eV}$, as is standard for ILs.^{51, 66, 67}

In some cases, calculated XP spectra were shifted in E_B to give the best visual match; E_B shifts used are given in the Figure captions. Calculated Zn 1s XANES spectra were shifted -8.9 eV in energy, which has been found to give excellent matches to experimental XANES spectra.⁶⁸

XPS of $[\text{C}_8\text{C}_1\text{Im}]\text{Cl}$, *i.e.* $x = 0.00$ with Cl^- solvated by $[\text{C}_8\text{C}_1\text{Im}]^+$, gives one experimental Cl electronic environment in the Cl 2p core XP spectrum, which manifests as two peaks due to spin-orbit coupling; $\Delta E_B(\text{Cl } 2p_{3/2} - \text{Cl } 2p_{1/2}) = 1.60\ \text{eV}$ and the peak area ratio for Cl $2p_{3/2}$ to Cl $2p_{1/2}$ is 2:1 (Figure 2a). The same occurs for $[\text{C}_8\text{C}_1\text{Im}]\text{Br}$, where XPS gives one experimental Br electronic environment in the Br 3d core XP spectrum (Figure 2c); $\Delta E_B(\text{Br } 3d_{5/2} - \text{Br } 3d_{3/2}) = 1.04\ \text{eV}$ and the peak area ratio for Br $3d_{5/2}$ to Br $3d_{3/2}$ is 3:2 (Figure 2c). On a macroscopic level, $[\text{C}_8\text{C}_1\text{Im}]\text{Cl}$ and $[\text{C}_8\text{C}_1\text{Im}]\text{Br}$ both present one halide electronic environment in the liquid phase with FWHM of $\sim 0.90\ \text{eV}$ and $\sim 0.85\ \text{eV}$ respectively (ESI Table S5). Therefore, we use the Cl 2p XP spectrum for $[\text{C}_8\text{C}_1\text{Im}]\text{Cl}$ and the Br 3d XP spectrum for $[\text{C}_8\text{C}_1\text{Im}]\text{Br}$ as our baseline for judging one halide electronic environment in the liquid phase.

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3. Results

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3.1. Speciation at $x = 0.33$: $1 \times \text{ZnHal}_2 + 2 \times \text{Hal}^-$ forms $[\text{ZnHal}_4]^{2-}$

For $x = 0.00$ and $x = 0.33$ the experimental and calculated Cl 2p match both in terms of E_B differences and FWHM (Figure 2a) which indicates that all four Cl are equivalent for $x = 0.33$. The core XPS makes clear that there are four terminal chlorides in each $[\text{ZnCl}_4]^{2-}$ anion, with no other anion species detectable, *i.e.* no Cl^- , $[\text{Zn}_2\text{Cl}_6]^{2-}$ or $[\text{ZnCl}_3]^-$. Therefore, anion speciation for $x = 0.33$ is $[\text{ZnCl}_4]^{2-}$. The same results are found for $x = 0.33$ with Br; the experimental and calculated Br 3d match for $x = 0.00$ and $x = 0.33$ both in terms of E_B differences and FWHM (Figure 2b). Therefore, anion speciation for $x = 0.33$ is $[\text{ZnBr}_4]^{2-}$. Our results match well to findings from Raman spectroscopy and EXAFS,^{16, 18, 37} and demonstrate that the gas phase $[\text{ZnCl}_3]^-$ speciation observed using mass spectrometry^{30, 32} does not match the liquid phase speciation.

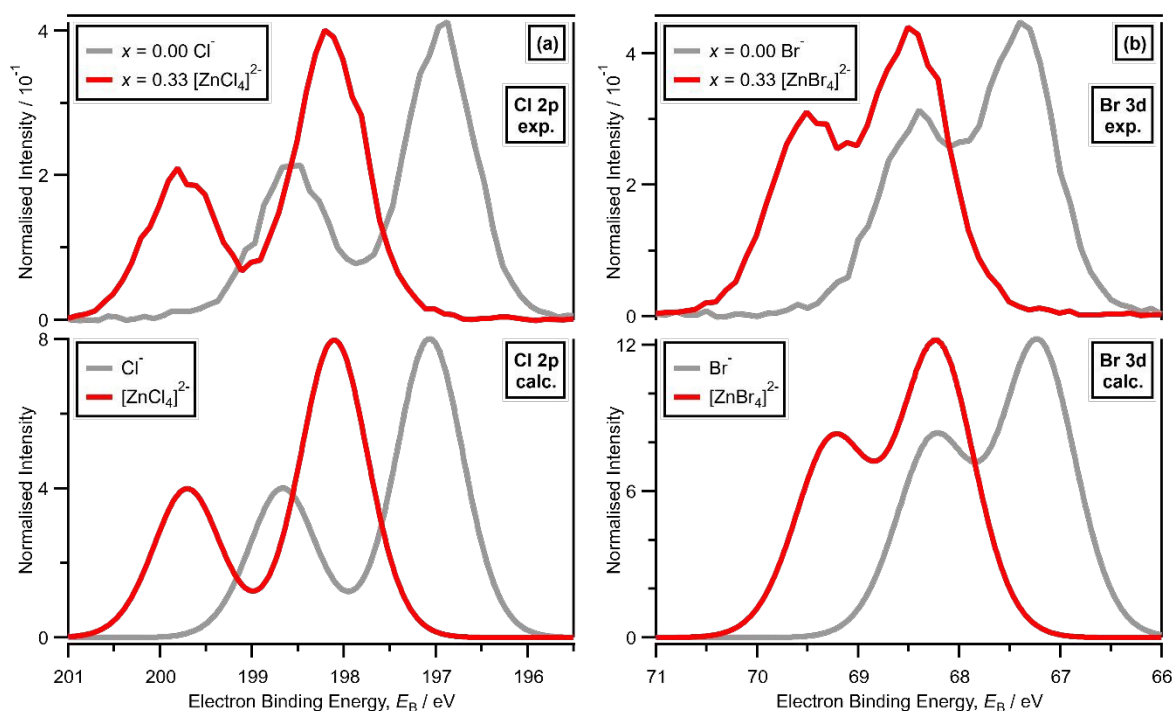


Figure 2. (a) (top) Experimental core Cl 2p XPS for $x = 0.00$ $[\text{C}_8\text{C}_1\text{Im}]\text{Cl}$ and $x = 0.33$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{ZnCl}_4]$ and (bottom) lone ion SMD calculated core Cl 2p XPS for Cl^- and $[\text{ZnCl}_4]^{2-}$ (both shifted by $E_B = 2.65$ eV). (b) (top) Experimental core Br 3d XPS for $x = 0.00$ $[\text{C}_8\text{C}_1\text{Im}]\text{Br}$ and $x = 0.33$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{ZnBr}_4]$ and (bottom) lone ion SMD calculated core Br 3d XPS for Br^- and $[\text{ZnBr}_4]^{2-}$ (both shifted by $E_B = -2.36$ eV).

3.2. Speciation at $x = 0.50$: $2 \times \text{ZnHal}_2 + 2 \times \text{Hal}^-$ forms $[\text{Zn}_2\text{Hal}_6]^{2-}$ and not $2 \times [\text{ZnHal}_3]^-$

For $x = 0.50$, *i.e.* $2 \times \text{ZnHal}_2 + 2 \times \text{Hal}^-$, more than one halide electronic environment exists for both Cl and Br, as the Cl 2p and Br 3d photoemission profiles for $x = 0.50$ do not visually match the shape of the Cl 2p and Br 3d XP spectra respectively (Figure 2), confirming that there is more than one halide environment for $x = 0.50$. Therefore, at $x = 0.50$ the speciation cannot be $[\text{ZnHal}_3]^-$ only (with no $[\text{Zn}_2\text{Cl}_6]^{2-}$), as $[\text{ZnHal}_3]^-$ would give a single halide environment in Cl 2p/Br 3d XPS. The visual match of experimental and calculated core and valence XPS for both Cl and Br for the experimental $x = 0.50$ versus calculated $[\text{Zn}_2\text{Hal}_6]^{2-}$ are almost perfect (Figure 3a to Figure 3d). The core XPS makes clear that there are two bridging and four terminal halides in each $[\text{Zn}_2\text{Hal}_6]^{2-}$ anion (which in an ideal structure has D_{2h} symmetry), matching the speciation found using Raman spectroscopy¹⁶. Furthermore, the visual match of experimental and calculated core and valence XPS for both Cl and Br for the experimental $x = 0.50$ versus $[\text{ZnHal}_3]^-$ are relatively poor (Figure 3a to Figure 3d). Moreover, the



experimental Zn 1s XANES spectroscopy for $x = 0.50$ for both Cl and Br match well to the calculated Zn 1s XANES spectroscopy for $[\text{Zn}_2\text{Hal}_6]^{2-}$ and very poorly to $[\text{ZnHal}_3]^-$ (Figure 3e and Figure 3f). Therefore, the $[\text{ZnCl}_3]^-$ speciation proposed from gas phase mass spectrometry results^{28, 30} does not reflect the liquid phase speciation. Overall, our results show definitively that $x = 0.50$ (*i.e.* $2 \times \text{ZnCl}_2 + 2 \times \text{Cl}$) gives $[\text{Zn}_2\text{Cl}_6]^{2-}$ and not $2 \times [\text{ZnCl}_3]^-$.

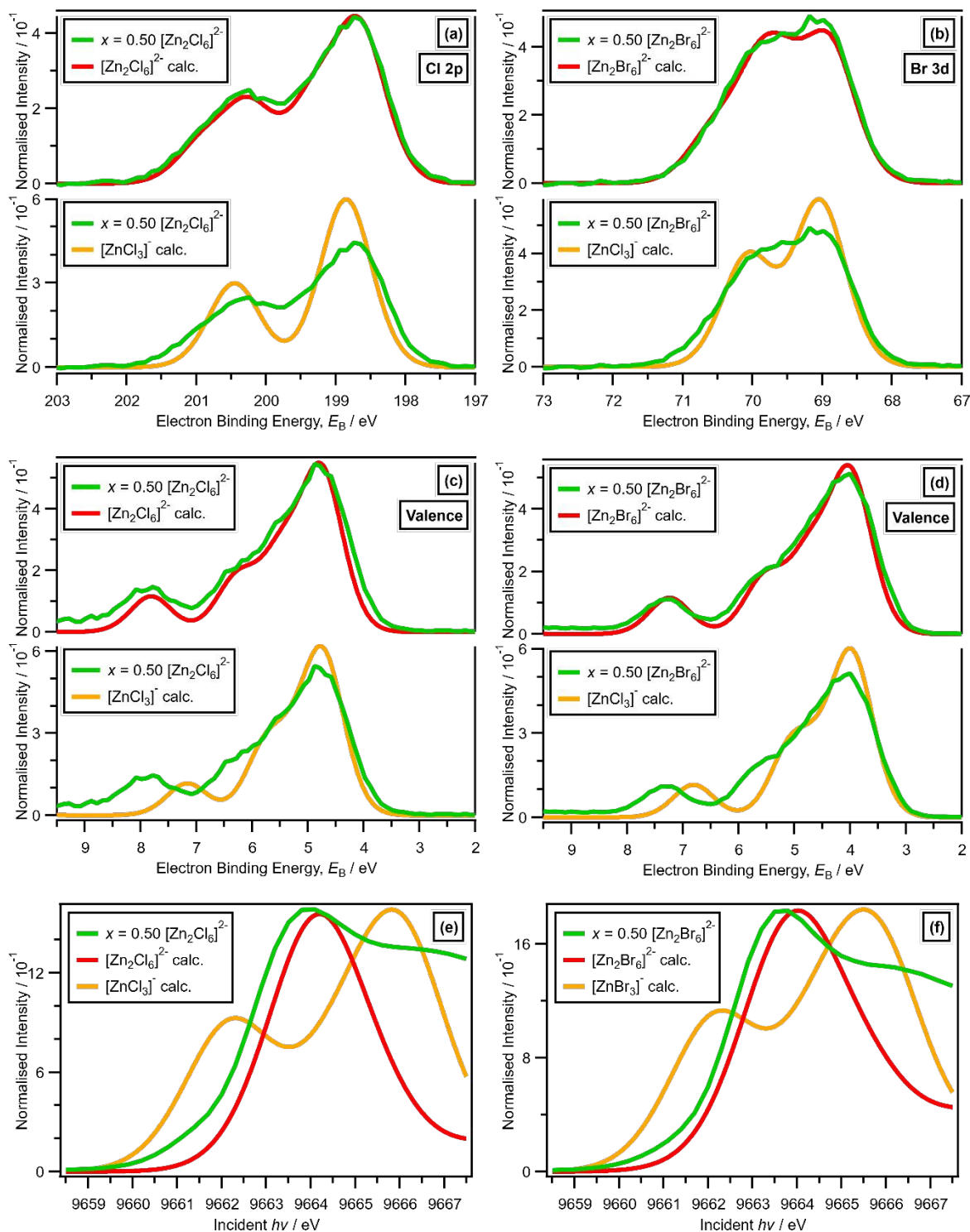


Figure 3. (a) Experimental core Cl 2p XPS for $x = 0.50$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_2\text{Cl}_6]$ and (top) lone ion SMD calculated core Cl 2p XPS for $[\text{Zn}_2\text{Cl}_6]^{2-}$ (shifted by $E_B = 2.65$ eV) and (bottom) lone ion SMD calculated core Cl 2p XPS $[\text{ZnCl}_3]^-$ (shifted by $E_B = 2.50$ eV). (b) Experimental core Br 3d XPS for $x = 0.50$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_2\text{Br}_6]$ and (top) lone ion SMD calculated core Br 3d XPS for $[\text{Zn}_2\text{Br}_6]^{2-}$ (shifted by $E_B = -2.36$ eV) and (bottom) lone ion SMD calculated core Br 3d XPS



[ZnBr₃]⁻ (shifted by $E_B = -2.82$ eV). (c) Experimental valence XPS for $x = 0.50$ [C₈C₁Im]₂[Zn₂Cl₆] and (top) lone ion SMD calculated valence XPS for [Zn₂Cl₆]²⁻ (shifted by $E_B = -4.47$ eV) and (bottom) lone ion SMD calculated valence XPS for [ZnCl₃]⁻ (shifted by $E_B = -4.77$ eV). (d) Experimental valence XPS for $x = 0.50$ [C₈C₁Im]₂[Zn₂Br₆] and (top) lone ion SMD calculated valence XPS for [Zn₂Br₆]²⁻ (shifted by $E_B = -4.23$ eV) and (bottom) lone ion SMD calculated valence XPS for [ZnBr₃]⁻ (shifted by $E_B = -4.77$ eV). (e) Experimental Zn 1s XANES for $x = 0.50$ [C₈C₁Im]₂[Zn₂Cl₆], and lone ion SMD calculated Zn 1s XANES for [Zn₂Cl₆]²⁻ and [ZnCl₃]⁻ (shifted by incident $h\nu = -8.9$ eV). (f) Experimental Zn 1s XANES for $x = 0.50$ [C₈C₁Im]₂[Zn₂Br₆], and lone ion SMD calculated Zn 1s XANES for [Zn₂Br₆]²⁻ and [ZnBr₃]⁻ (shifted by incident $h\nu = -8.9$ eV).

3.3. Speciation at $x = 0.67$: $4 \times \text{ZnHal}_2 + 2 \times \text{Hal}^-$ forms linear and supertetrahedron [Zn₄Hal₁₀]²⁻

$x = 0.67$, *i.e.* $2 \times \text{ZnHal}_2 + 1 \times \text{Hal}^-$, gave more than one halide electronic environment, as the Cl 2p (and Br 3d) for $x = 0.67$ does not match the Cl 2p (and Br 3d) for $x = 0.00$ (Figure 6a and Figure 7a). Furthermore, the visual match of experimental and calculated core and valence XPS for both Cl and Br for the experimental $x = 0.67$ versus [Zn₄Hal₁₀]²⁻ (both linear with D_{2h} symmetry and supertetrahedron with T_d symmetry, Figure 1) are good (Figure 4, Figure 6 and Figure 7), matching a finding from Raman spectroscopy of linear [Zn₄Cl₁₀]²⁻.¹⁶ The core XPS makes clear that there are six bridging and four terminal halides in each [Zn₄Hal₁₀]²⁻ anion (ESI Figure S7d and Figure S12d). Therefore, a structure that has been suggested in the literature,³⁶ [Zn₂Cl₅]⁻ with three bridging Cl between the two Zn atoms and one terminal Cl on each Zn atom (ESI Figure S1), is certainly not present, confirmed by the visual mismatch between experimental and calculated XP spectra (both core Cl 2p and valence, ESI Figure S22 and Figure S23). Furthermore, [Zn₂Cl₅]⁻ with two bridging Cl between the two Zn atoms, two terminal Cl on one Zn atom, and one terminal Cl on one Zn atom could not be optimised using DFT calculations. However, the core XPS is not suitable for distinguishing linear and supertetrahedron contributions. The visual match of experimental and calculated valence XPS for both Cl and Br for the experimental $x = 0.67$ versus [Zn₄Hal₁₀]²⁻ are good (Figure 4 top and middle). However, a 1:1 combination of calculated linear and supertetrahedron valence XPS gives an even better visual match (Figure 4 bottom) than the individual linear and supertetrahedron valence XPS (Figure 4 top and middle). The uncertainty in 1:1 ratio is relatively large, but we are confident that the speciation for $x = 0.67$ is not only one species. Overall, we found good evidence that two species are present for $x = 0.67$, linear and supertetrahedron [Zn₄Cl₁₀]²⁻, in approximately 1:1 ratio, with no evidence for the presence of 1- anions such as [ZnCl₃]⁻, [Zn₂Cl₅]⁻, [Zn₃Cl₇]⁻ observed using mass spectrometry and found from EXAFS data.^{28-30, 32}



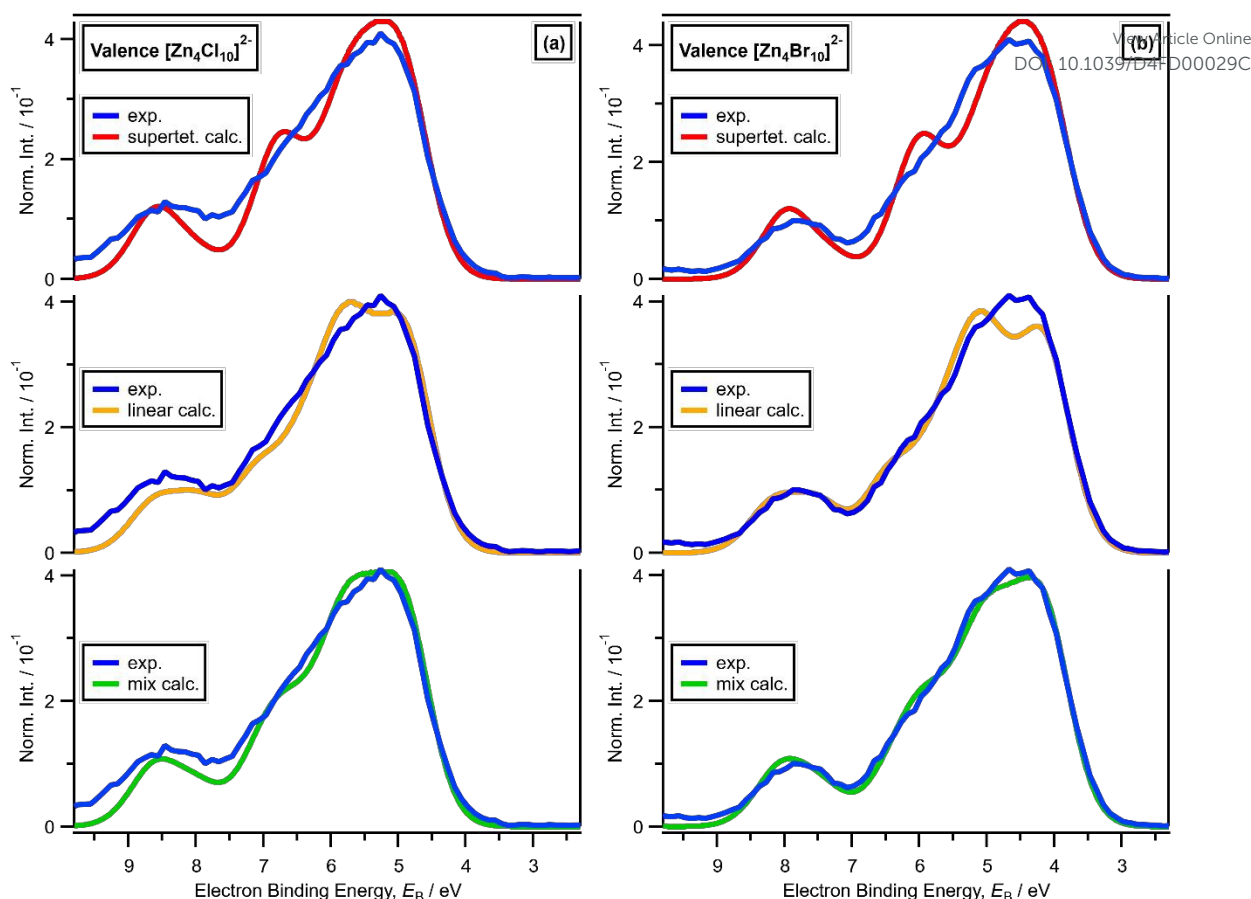


Figure 4. (a) Experimental valence XPS for $x = 0.67$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_4\text{Cl}_{10}]$ and (top) lone ion SMD calculated valence XPS for supertetrahedron $[\text{Zn}_4\text{Cl}_{10}]^{2-}$, (middle) linear $[\text{Zn}_4\text{Cl}_{10}]^{2-}$, and (bottom) 1:1 mix of supertetrahedron $[\text{Zn}_4\text{Cl}_{10}]^{2-}$ and linear $[\text{Zn}_4\text{Cl}_{10}]^{2-}$ (shifted by $E_B = -4.67$ eV). (b) Experimental valence XPS for $x = 0.67$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_4\text{Br}_{10}]$ and (top) lone ion SMD calculated valence XPS for supertetrahedron $[\text{Zn}_4\text{Br}_{10}]^{2-}$, (middle) linear $[\text{Zn}_4\text{Br}_{10}]^{2-}$, and (bottom) 1:1 mix of supertetrahedron $[\text{Zn}_4\text{Br}_{10}]^{2-}$ and linear $[\text{Zn}_4\text{Br}_{10}]^{2-}$ (shifted by $E_B = -4.55$ eV).

3.4. Speciation at $x = 0.33, 0.50, 0.60$ for halozincates: one anion present

Results from Section 3.1 for $x = 0.33$ and Section 3.2 for $x = 0.50$ all point towards there being only one species present, $[\text{ZnCl}_4]^{2-}$ and $[\text{Zn}_2\text{Cl}_6]^{2-}$ respectively. Results from Section 3.3 show that there were two anions present for $x = 0.67$, linear and supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$. For $x = 0.43$, it is impossible for there to be only one anion present; two anions must be present, predicted to be a 1:1 ratio of $[\text{ZnCl}_4]^{2-}$ and $[\text{Zn}_2\text{Cl}_6]^{2-}$.

A larger FWHM($\text{Zn } 2p_{3/2}$) for $x = 0.43$ was clearly observed, relative to $x = 0.33$ and $x = 0.50$ (with Cl ligands), when the Zn $2p_{3/2}$ XP spectra are set at the same $E_B(\text{Zn } 2p_{3/2})$ and normalised to the same peak intensity to make visual judgements of the FWHM relatively easy (Figure 5a). FWHM($\text{Zn } 2p_{3/2}$) from peak fitting (Figure 5b) gave the same results as the visual assessment of the peak FWHM (Figure 5a). The order of FWHM($\text{Zn } 2p_{3/2}$) (and FWHM($\text{Zn } 2p_{1/2}$), ESI Figure S24) for x was: $0.33 < 0.50 < 0.60 \approx 0.67 < 0.43$ (Figure 5b). Therefore, FWHM($\text{Zn } 2p_{3/2}$) was far larger for $x = 0.43$ when two species were definitely present than all other x (Figure 5b). For $x = 0.43$ and $x = 0.67$, two anions were present for both solutions, but FWHM($\text{Zn } 2p_{3/2}$) was substantially larger for $x = 0.43$ than $x = 0.67$, because the two anions present for $x = 0.67$ (linear and supertetrahedron $[\text{Zn}_4\text{Cl}_{10}]^{2-}$) were more similar than the two anions present for $x = 0.43$ ($[\text{ZnCl}_4]^{2-}$ and $[\text{Zn}_2\text{Cl}_6]^{2-}$).



There is a very good match of experimental and calculated XPS for both $E_B(\text{Cl } 2p)$ and XP spectral shape as x increased for Cl-containing ILs (Figure 6); the same observations hold for $E_B(\text{Br } 3d)$ as x increased for Br-containing ILs (Figure 7). These findings provide more evidence that one halozincate anion exists at certain x .

These observations confirm that one halozincate anion was present for $x = 0.33$, $x = 0.50$ and $x = 0.60$, which are $[\text{ZnHal}_4]^{2-}$, $[\text{Zn}_2\text{Hal}_6]^{2-}$ and $[\text{Zn}_3\text{Hal}_8]^{2-}$ respectively (Figure 8a). Furthermore, at $x = 0.67$ only $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ is present, most likely a 1:1 or similar combination of linear and supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$. This finding contrasts with results for $\text{AlCl}_3 + \text{Cl}^-$,¹ where multiple anions were proposed where there was the potential to form only one anion, *e.g.* $x = 0.67$ (Figure 8b).

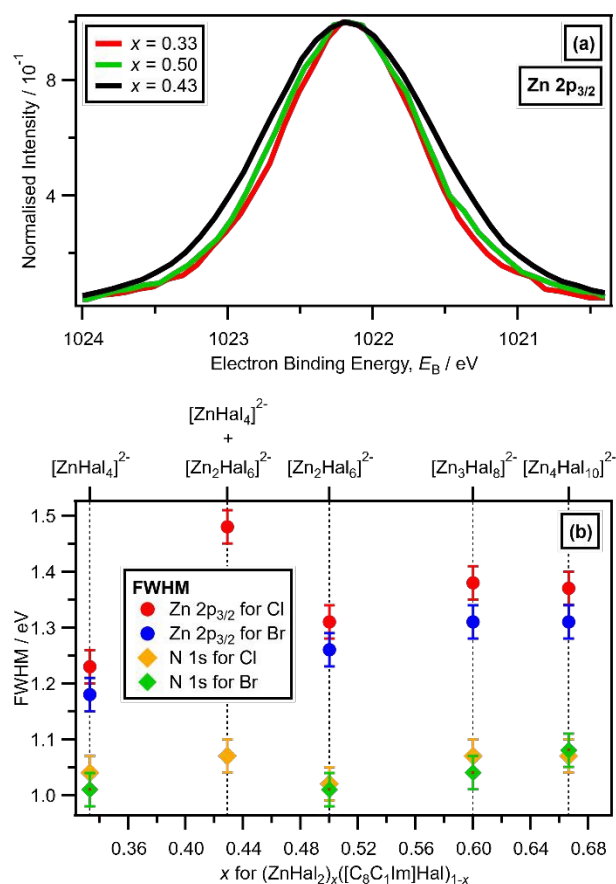


Figure 5. (a) Experimental Zn $2p_{3/2}$ XPS for $x = 0.33$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{ZnCl}_4]$, for $x = 0.50$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_2\text{Cl}_6]$, and for $x = 0.43$ $[\text{C}_8\text{C}_1\text{Im}]_4[\text{ZnCl}_4][\text{Zn}_2\text{Cl}_6]$ (peak intensity normalised to 1, shifted so all $E_B(\text{Zn } 2p_{3/2}) = 1022.16$ eV). (b) Experimental XPS FWHM for Zn $2p_{3/2}$, Zn $2p_{1/2}$, Zn $2p_{3/2}$ and N_{cation} 1s for different x for $([\text{C}_8\text{C}_1\text{Im}]\text{Hal})_{1-x}(\text{ZnHal}_2)_x$ from $x = 0.33$ to $x = 0.67$.



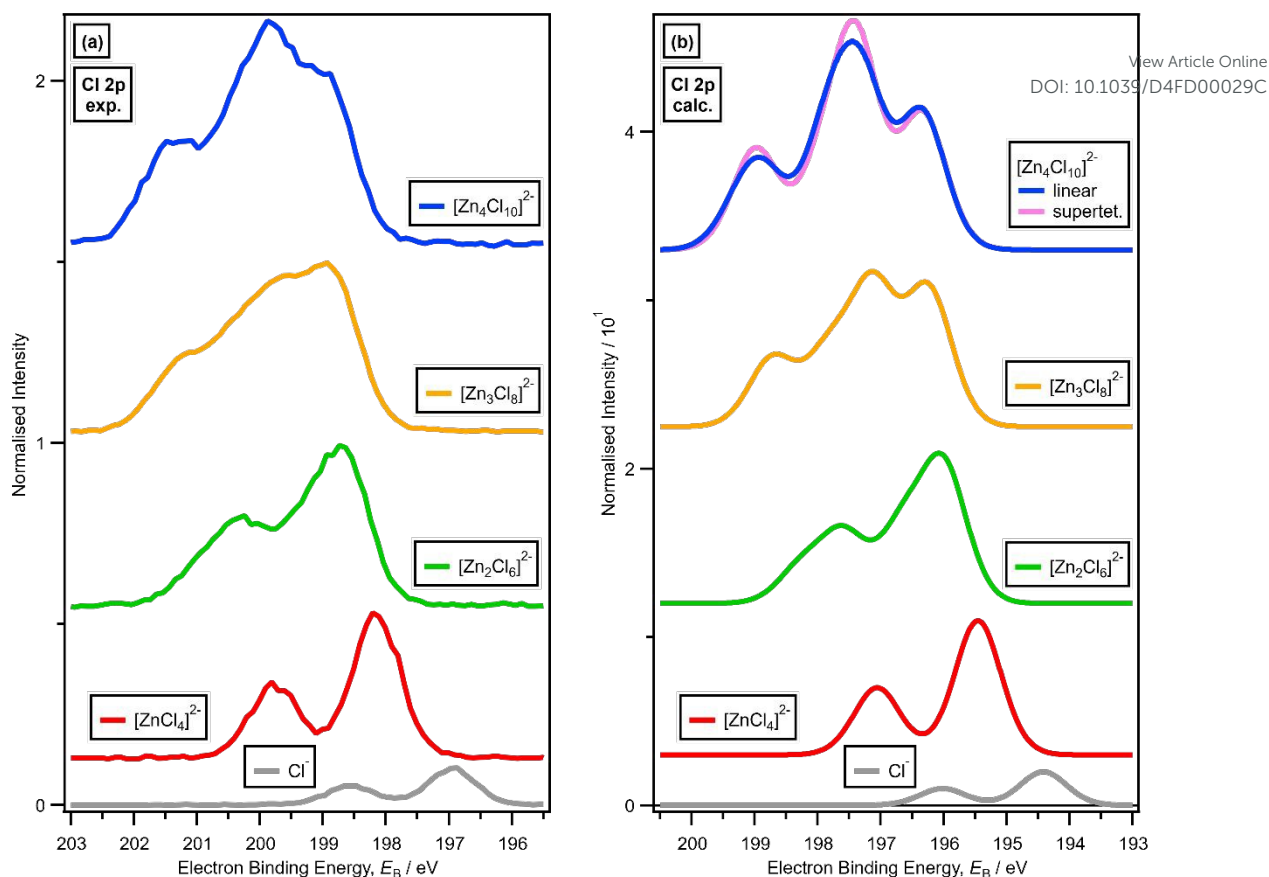


Figure 6. (a) Experimental core Cl 2p XPS for $x = 0.00$ $[\text{C}_8\text{C}_1\text{Im}]\text{Cl}$, $x = 0.33$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{ZnCl}_4]$, $x = 0.50$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_2\text{Cl}_6]$, $x = 0.60$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_3\text{Cl}_8]$, $x = 0.67$ $[\text{C}_8\text{C}_1\text{Im}]_2[\text{Zn}_4\text{Cl}_{10}]$ (vertically offset for clarity). (b) Lone ion SMD calculated core Cl 2p XPS for Cl^- , $[\text{ZnCl}_4]^{2-}$, $[\text{Zn}_2\text{Cl}_6]^{2-}$, $[\text{Zn}_3\text{Cl}_8]^{2-}$ and $[\text{Zn}_4\text{Cl}_{10}]^{2-}$ (vertically offset for clarity).



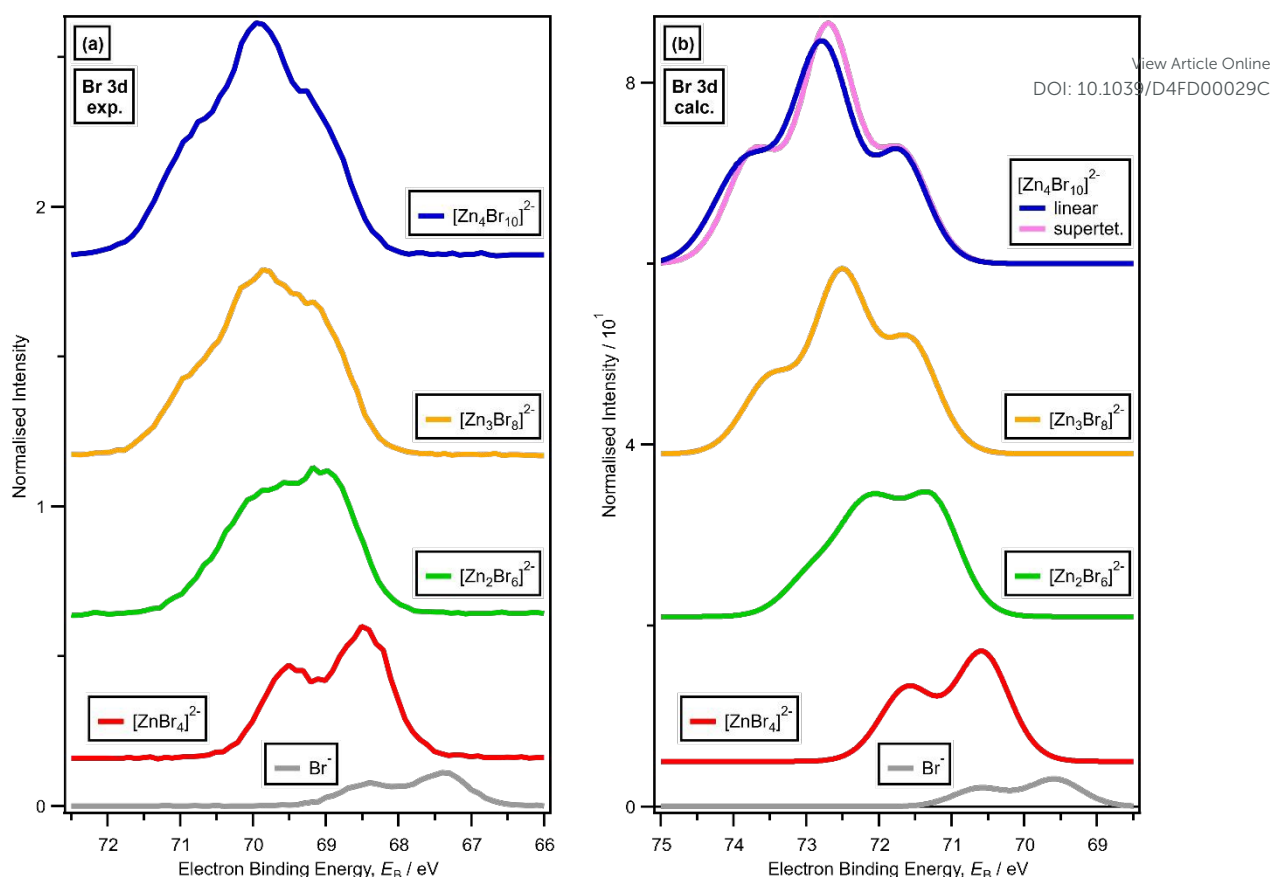


Figure 7. (a) Experimental core Br 3d XPS for $x = 0.00$ [C_8C_1Im]Br, $x = 0.33$ [C_8C_1Im] $_2$ [ZnBr $_4$] $^{2-}$, $x = 0.50$ [C_8C_1Im] $_2$ [Zn $_2$ Br $_6$] $^{2-}$, $x = 0.60$ [C_8C_1Im] $_2$ [Zn $_3$ Br $_8$] $^{2-}$, $x = 0.67$ [C_8C_1Im] $_2$ [Zn $_4$ Br $_{10}$] $^{2-}$ (vertically offset for clarity). (b) Lone ion SMD calculated core Br 3d XPS for Br $^-$, [ZnBr $_4$] $^{2-}$, [Zn $_2$ Br $_6$] $^{2-}$, [Zn $_3$ Br $_8$] $^{2-}$ and [Zn $_4$ Br $_{10}$] $^{2-}$ (vertically offset for clarity).

3.5. Chlorozincate and bromozincate speciation match at all x

All data presented so far in this article (Figure 2 to Figure 7) shows that chlorozincate and bromozincate ILs have the same anion speciation at any x . One last piece of evidence is that both the experimental and calculated valence XPS for each x studied have the same spectral features, *i.e.* the anion(s) present for each x has the same bonding and therefore the same speciation whether the ligands are Cl or Br (ESI Figure S25). For example, at $x = 0.33$ there are clearly three groups of features for both chlorozincate and bromozincate (ESI Figure S25a and S25d), *i.e.* the bonding and therefore the speciation was the same, [ZnHal $_4$] $^{2-}$. The same feature comes at $E_B \sim 0.7$ eV lower in the experimental valence XPS for bromozincate anions compared to the chlorozincate anions (ESI Figure S25a and S25c), and at $E_B \sim 1.0$ eV lower in the calculated valence XPS for bromozincate anions compared to the chlorozincate anions (ESI Figure S25d and S25f).

3.6. Halozincate speciation: discussion

All our experimental evidence, backed up by calculations, shows that only 2- halozincate anions occur in the liquid phase up to $x = 0.67$ (Figure 8a). For certain x only one anion formed within our detection limit (estimated to be 1% of the main species); $x = 0.33$ ([ZnHal $_4$] $^{2-}$), $x = 0.50$ ([Zn $_2$ Hal $_6$] $^{2-}$) and $x = 0.60$ ([Zn $_3$ Hal $_8$] $^{2-}$) (Figure 8a), demonstrating that any equilibria with other anions is very much to towards the halozincate anions listed here. Our liquid phase findings using X-ray spectroscopy contrast strongly to findings from gas phase mass spectrometry, where equilibria of multiple halozincate anions were observed²⁹. Clearly, mass spectrometry is an inadequate technique for studying speciation of halozincate anions in ILs,²² and probably for all anions with a formal negative charge of 2- or larger.



For $x = 0.67$, only one anion composition was found but with two different isomers, supertetrahedron and linear $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ (Figure 8a), suggesting that an equilibrium occurred for $x = 0.67$. We expect that the same trend will continue for larger x as ZnHal_2 do not form stable neutral dimers (unlike AlCl_3), e.g. at $x = 0.714$ we predict only $[\text{Zn}_5\text{Cl}_{12}]^{2-}$ would form. Overall, there are specific turning points in the halozincate speciation.

For all x , at least up to $x = 0.60$, there are definitely two 2- halozincate anions present for all non-turning point values of x (i.e. all x values apart from $x = 0.33$, $x = 0.50$ and $x = 0.60$, Figure 8a). It is tough to use X-ray spectroscopy to judge the exact halozincate speciation for x away from the turning point x values, as demonstrated by the Cl 2p XPS data for $x = 0.43$ (ESI Section 2). However, given the fact we have these turning point x values, we expect that at $0.33 < x < 0.67$ the speciation can be predicted to be a combination of the expected ratios of halozincate anions (Figure 8a). For example, $x = 0.556$ would give $1 \times [\text{Zn}_2\text{Cl}_6]^{2-} + 1 \times [\text{Zn}_3\text{Cl}_8]^{2-}$, $x = 0.538$ would give $2 \times [\text{Zn}_2\text{Cl}_6]^{2-} + 1 \times [\text{Zn}_3\text{Cl}_8]^{2-}$ and $x = 0.571$ would give $1 \times [\text{Zn}_2\text{Cl}_6]^{2-} + 2 \times [\text{Zn}_3\text{Cl}_8]^{2-}$ (Figure 8a).

IL Lewis acidity (measured using the Gutmann-Beckett method) has a step-change at $x = 0.33$,¹⁶ which matches to the change in speciation above $x = 0.33$ with the appearance of $[\text{Zn}_2\text{Cl}_6]^{2-}$. Furthermore, the Lewis acidity shows no significant step changes with relatively gentle increases when x was increased to above $x = 0.50$ or above $x = 0.60$,¹⁶ even though we know new halozincate anions appear above those x values. Therefore, the presence of bridging Zn-Hal bonds appears to be essential for high halozincate Lewis acidity. Zinc centres with four halide bonds do not appear capable of increasing their coordination number to allow the bonding of the Lewis basic probe, triethylphosphine oxide (TEPO), as $[\text{ZnCl}_4(\text{OH}_2)]^{2-}$ and $[\text{ZnBr}_4(\text{OH}_2)]^{2-}$ are calculated to be thermodynamically unstable species.⁷⁰⁻⁷² Therefore, for halozincate anions the TEPO probe would have to bind to a zinc centre after the breaking of a Zn-Hal bond. We propose that bridging Zn-Hal bonds dissociate more readily than terminal Zn-Hal bonds, meaning that halozincate anions with bridging Zn-Hal bonds have higher Lewis acidity than $[\text{ZnHal}_4]^{2-}$. The smaller changes in Lewis acidity as x increased above 0.33 were likely due to a combination of bridging Zn-Hal bonds being easier to dissociate for larger halozincate anions (apart from supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ we expect) and changes in the electronic structure.

Thermal stability increases with increasing x ,¹⁷ which can be related principally to metal complex speciation. It has been proposed that the presence of free Hal^- is required for thermal decomposition of halozincate ILs.¹⁷ Hence, thermal stability shows a significant increase from $x = 0.00$ to $x = 0.33$, where only $[\text{ZnCl}_4]^{2-}$ and no free Hal^- is present.¹⁷ Thermal stability continues to increase with increasing x beyond $x = 0.33$, indicating that the presence of halozincate anions with bridging Hal leads to increased thermal stability, likely due to the greater energy barrier to forming free Hal^- anions.

The viscosity of halozincate ionic liquids is larger than their haloaluminate equivalents.²⁹ This greater viscosity is very likely linked to the speciation, i.e. the differences in charge of the anions $[\text{Zn}_n\text{Hal}_{2n+2}]^{2-}$ and $[\text{Al}_n\text{Hal}_{3n+1}]^-$. In general, ILs with 2- halometallate anions have stronger electrostatic interactions with organic cations than 1- halometallate anions, although that is not true across all halometallate anions.^{18, 19}

To fully understand many of the properties of halometallate-containing ILs, e.g. anion redox properties, both the speciation and the electronic structure need to be considered. The electronic structure of halozincate anions will be considered in a separate publication.⁷³

For some applications, e.g. catalysis, trace amounts of an unexpected halozincate anion could make data interpretation very challenging indeed. The chances of making an IL with only one x is expected to be extremely challenging. Commercial ZnCl_2 often contains traces of hydroxide, even those sold as anhydrous,⁷⁴ meaning that the composition a researcher intends to make may not quite occur.



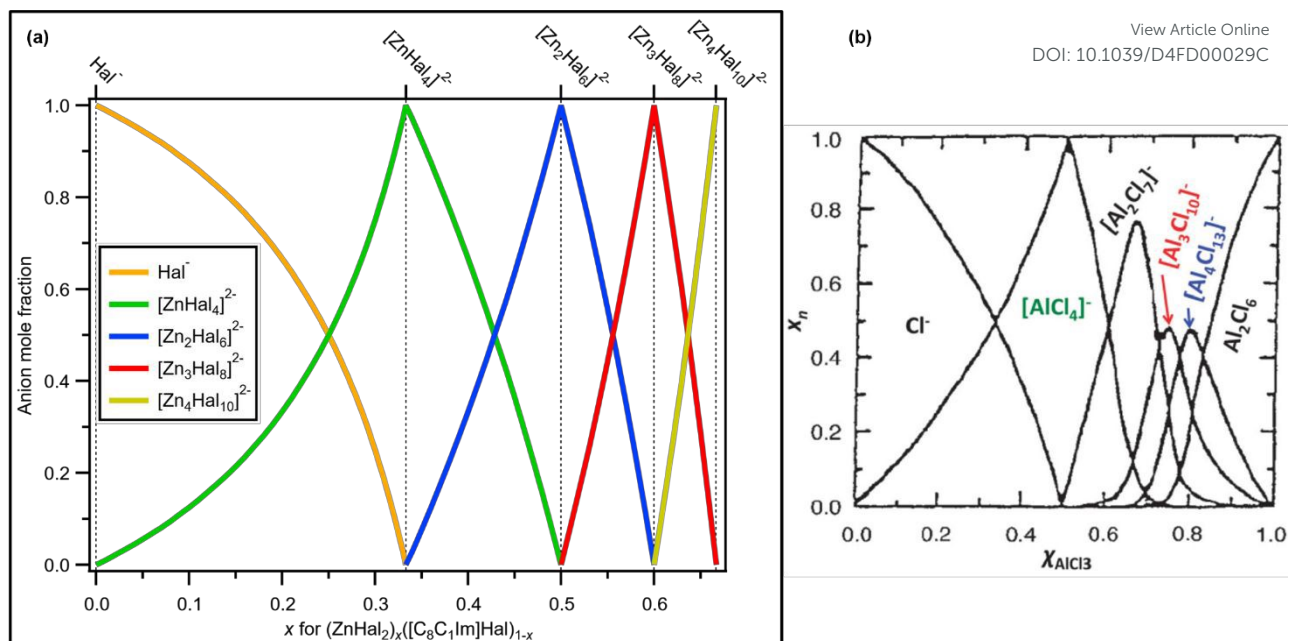


Figure 8. (a) Mole fraction of anions products plotted against mole fraction of zinc halide starting material for mixtures of ZnHal_2 and $[\text{C}_8\text{C}_1\text{Im}]\text{Hal}$. Values for $x = 0.00$, $x = 0.33$, $x = 0.50$, $x = 0.60$ and $x = 0.67$ are known; anion product speciation at other x is calculated. (b) Mole fraction of anions products plotted against mole fraction of aluminium chloride starting material for mixtures of AlCl_3 and $[\text{C}_2\text{C}_1\text{Im}]\text{Hal}$ (at 200°C , calculated from a thermodynamic model; taken with permission from reference ¹, original data from reference ⁶⁹).

3.7. Attempting to use differences in calculated total energies to predict speciation

The introduction of an SMD provides an adequate description/approximation of the surrounding solvent environment without actually explicitly including cations and additional anions. This is a big bonus, especially as it allows for the use of large basis sets (as in this case).

In the gas phase, total energies from calculations strongly suggest that 1- anions are massively more stable than 2- anions (ESI Table S6). For example, $2 \times [\text{ZnCl}_3]^- + 2 \times \text{Cl}^-$ are far more stable in the gas phase than $2 \times [\text{ZnCl}_4]^{2-}$ (ESI Table S6). Firstly, this finding is in complete contrast to our liquid phase experimental speciation. Secondly, this finding also strongly indicates why anion speciation from mass spectrometry techniques is unlikely to match liquid phase speciation, as gas phase stability must be a strong factor in the halozincate anions observed in the mass spectrometry.

For calculations using the SMD, there are a number of examples where the halozincate anion speciation predicted by the calculated total energies does not match our experimental findings according to ΔG (Table 1): (i) for $x = 0.33$, $1 \times [\text{ZnCl}_3]^- + 1 \times \text{Cl}^-$ is considerably more stable than $2 \times [\text{ZnCl}_4]^{2-}$ and $1 \times [\text{ZnBr}_3]^- + 1 \times \text{Br}^-$ is the same stability as $2 \times [\text{ZnBr}_4]^{2-}$; (ii) for $x = 0.50$, $2 \times [\text{ZnHal}_3]^-$ is considerably more stable than $1 \times [\text{Zn}_2\text{Hal}_6]^{2-}$ for both Cl and Br ligands; (iii) for $x = 0.67$, $1 \times$ supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ is considerably more stable than $1 \times$ linear $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ for both Cl and Br ligands; (iv) for $x = 0.60$, $1 \times [\text{Zn}_2\text{Hal}_6]^{2-} + 1 \times$ supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ is more stable than $2 \times [\text{Zn}_3\text{Hal}_8]^{2-}$. Of these mismatches of experimental data and calculated total energy differences, numbers (i), (ii) and (iv) in our list above are the ones that make us doubt the reliability of using calculated total energy differences to judge anion speciation, as we have very high confidence in our experimental measurements of anion speciation for $x = 0.33$, $x = 0.50$ and $x = 0.60$. For mismatch (iii) listed above, we are less confident of our experimental measurements of anion speciation for $x = 0.67$.



It is clear from the calculations that by total energy differences the supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ is judged as far more stable than the linear $[\text{Zn}_4\text{Hal}_{10}]^{2-}$. It is possible that the calculated total energy differences are simply unreliable in this case. It could also be that kinetic contributions are important here, that the supertetrahedron $[\text{Zn}_4\text{Hal}_{10}]^{2-}$ takes a relatively large amount of energy to form. Such discussions are beyond the scope of our calculations.

Two different energy values (at least) have been used for judging halozincate speciation: Gibbs free energies, ΔG , and sum of electronic and zero-point energies, ΔE (Table 1). If only total energy differences were used to judge speciation, different conclusions would be drawn if either ΔG or ΔE were used (Table 1). For example, for $x = 0.33$ ΔG predicts $2 \times [\text{ZnCl}_3]^- + 2 \times \text{Cl}^-$ is more stable than $2 \times [\text{ZnCl}_4]^{2-}$, whereas the reverse is true using ΔE (Table 1) and also our conclusions from experimental data. Overall, it is clear that ΔE is better for judging speciation than ΔG (Table 1), as ΔE predicts speciation that matches experimental data more often than ΔG . However, neither ΔE nor ΔG match experimental speciation at all x (Table 1), demonstrating that small-scale DFT calculation total energy differences should be treated with caution when judging speciation of halozincate anions, and most likely all halometallate anions.

There are two reasons for this mismatch of speciation from experimental and calculated total energy differences. Firstly, the relative permittivity we are using is not correct, it is likely too low. We expect total energy differences from ΔG values will be very sensitive to the solvation environment, in our case the choice of the relative permittivity used for the SMD. We estimate that the relative permittivity needs to be set to ~ 40 (rather than the value of 11.4 that we have used as standard here) for ΔG to predict $x = 0.33$ that $2 \times [\text{ZnCl}_4]^{2-}$ is more stable than $2 \times [\text{ZnCl}_3]^- + 2 \times \text{Cl}^-$ (ESI Table S6). Secondly, the local structures around the halozincate anions are important for energetics. When using lone anions in an SMD, specific anion-cation interactions are clearly not included in calculations, and these interactions may be important for capturing true total energy differences. A major drawback for capturing these specific anion-cation interactions is at least two cations would be required for each 2-halozincate anion to achieve charge neutrality, and even larger clusters may be needed, which would be prohibitively expensive, especially with the DFT methods used in this article.

Overall, we have demonstrated that anion total energies from small-scale DFT calculations (used for example for halobismuthate anions²⁴) are not suitable as a standalone method for determining liquid phase halozincate speciation. Experimental data, ideally from both core and valence XPS and at times backed up by XANES spectroscopy, is required to have high confidence in judging anion speciation for halometallate ILs.

Attempts have been made to use calculated total energies (which can be used to calculate dissociation energies) to judge halozincate anion thermal stability.¹⁷ Our work suggests that ΔE and not ΔG would be far better placed to judge thermal stability.



Table 1. Experimental and calculated (using an SMD) speciation for chlorozincate and bromozincate anions The red text in columns 4 and 5 shows calculations that do not agree with the experimental speciation

ZnHal ₂ mole fraction, x	Reactants	Possible anionic species ^a	ΔE / kJ mol ⁻¹	ΔG / kJ mol ⁻¹
0.33	4 × Cl ⁻ + 2 × ZnCl ₂	2 × [ZnCl₄]²⁻	45	35
		2 × Cl ⁻ + 1 × [Zn ₂ Cl ₆] ²⁻ 2 × Cl ⁻ + 2 × [ZnCl ₃] ⁻	45	-24
0.33	4 × Br ⁻ + 2 × ZnBr ₂	2 × [ZnBr₄]²⁻	31	32
		2 × Br ⁻ + 1 × [Zn ₂ Br ₆] ²⁻ 2 × Br ⁻ + 2 × [ZnBr ₃] ⁻	57	4
0.43	4 × Cl ⁻ + 3 × ZnCl ₂	1 × [ZnCl₄]²⁻ + 1 × [Zn₂Cl₆]²⁻	22	-12
		1 × Cl ⁻ + 1 × [ZnCl ₃] ⁻ + 1 × [Zn ₂ Cl ₆] ²⁻ 1 × Cl ⁻ + 3 × [ZnCl ₃] ⁻	22	-71
0.50	4 × Cl ⁻ + 4 × ZnCl ₂	2 × [Zn₂Cl₆]²⁻	33	32
		1 × [ZnCl ₄] ²⁻ + 1 × [Zn ₃ Cl ₈] ²⁻ 4 × [ZnCl ₃] ⁻	0	-118
0.50	4 × Br ⁻ + 4 × ZnBr ₂	2 × [Zn₂Br₆]²⁻	36	27
		1 × [ZnBr ₄] ²⁻ + 1 × [Zn ₃ Br ₈] ²⁻ 4 × [ZnBr ₃] ⁻	53	-57
0.60	4 × Cl ⁻ + 6 × ZnCl ₂	2 × [Zn₃Cl₈]²⁻	13	21
		1 × [Zn ₂ Cl ₆] ²⁻ + 1 × linear [Zn ₄ Cl ₁₀] ²⁻ 1 × [Zn ₂ Cl ₆] ²⁻ + 1 × supertetrahedron [Zn ₄ Cl ₁₀] ²⁻	-30	-17
0.60	4 × Br ⁻ + 6 × ZnBr ₂	2 × [Zn₃Br₈]²⁻	17	18
		1 × [Zn ₂ Br ₆] ²⁻ + 1 × linear [Zn ₄ Br ₁₀] ²⁻ 1 × [Zn ₂ Br ₆] ²⁻ + 1 × supertetrahedron [Zn ₄ Br ₁₀] ²⁻	-25	-20
0.67	2 × Cl ⁻ + 4 × ZnCl ₂	1 × linear [Zn₄Cl₁₀]²⁻	-44	-38
		1 × supertetrahedron [Zn ₄ Cl ₁₀] ²⁻ 2 × [Zn ₂ Cl ₅] ⁻	61	10
0.67	2 × Br ⁻ + 4 × ZnBr ₂	1 × linear [Zn₄Br₁₀]²⁻	-41	-38
		1 × supertetrahedron [Zn ₄ Br ₁₀] ²⁻		

^a Experimental speciation given in bold in this column

4. Conclusions

We have made a comprehensive study of the speciation of halozincate anions in ILs using a combination of core XPS, valence XPS, Zn 1s XANES spectroscopy, DFT calculations for XPS and total energies, and time-dependent DFT calculations for XANES spectroscopy. We believe this is the first study that considers both 1- and 2- halozincate anions from a computational perspective. We have demonstrated that at specific x only one halozincate anion was present, and equilibria of different halozincate anions at those x do not occur to any significant level. Furthermore, we have made predictions of the speciation for all x up to 0.67. Small-scale DFT calculations were used in conjunction with experimental X-ray spectroscopy data to capture halozincate speciation, but we advise caution when using differences in calculated total energies to judge halozincate speciation. Lastly, we demonstrated that speciation for chlorozincates and bromozincates were the same at any particular x .

Our findings prove that high quality data from multiple spectroscopies and calculations are essential to have high confidence in halozincate anion speciation identification. Moreover, our methodology can be used to identify the speciation of any halometallate anion, which is especially important for the many spectroscopically quiet closed-shell metals, *e.g.* Ga, In, Bi. Furthermore, our methodology of using core XPS peak FWHM to judge speciation will be suitable for halometallates that are capable of expanding their coordination number, *e.g.* In (we have shown here (ESI Figure S26 and Figure S27) that $1 \times \text{InCl}_3 + 1 \times \text{Cl}^-$ forms exclusively (at our detection limit) $[\text{InCl}_4]^-$ as there is one Cl electronic environment, the same as Cl^-). ILs with more than one metal cation present have been shown to have complex speciation,⁷⁵ which our methods will also be suitable for. Lastly, catalysis using halometallate ILs is often run above room temperature; there are great challenges determining speciation in such cases.

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6. References

- J. Estager, J. D. Holbrey and M. Swadźba-Kwaśny, *Chem. Soc. Rev.*, 2014, **43**, 847-886.
- K. Li, H. Choudhary and R. D. Rogers, *Curr. Opin. Green Sustain. Chem.*, 2018, **11**, 15-21.
- A. P. Abbott and K. J. McKenzie, *Phys. Chem. Chem. Phys.*, 2006, **8**, 4265-4279.
- Z. Liu, T. Cui, T. Q. Lu, M. S. Ghazvini and F. Endres, *J. Phys. Chem. C*, 2016, **120**, 20224-20231.
- A. P. Abbott, G. Frisch and K. S. Ryder, in *Annual Review of Materials Research, Vol 43*, ed. D. R. Clarke, Annual Reviews, Palo Alto, 2013, vol. 43, pp. 335-358.
- C. Zhang, J. Holoubek, X. Y. Wu, A. Daniyar, L. D. Zhu, C. Chen, D. P. Leonard, I. A. Rodriguez-Perez, J. X. Jiang, C. Fang and X. Ji, *Chem. Commun.*, 2018, **54**, 14097-14099.
- X. Y. Wu, Y. K. Xu, C. Zhang, D. P. Leonard, A. Markir, J. Lu and X. L. Ji, *J. Am. Chem. Soc.*, 2019, **141**, 6338-6344.
- T. S. Zhang, Y. Tang, S. Guo, X. X. Cao, A. Q. Pan, G. Z. Fang, J. Zhou and S. Q. Liang, *Energy Environ. Sci.*, 2020, **13**, 4625-4665.
- X. C. Chen, D. D. Song, C. Asumana and G. R. Yu, *J. Mol. Catal. A-Chem.*, 2012, **359**, 8-13.
- L. H. Zhang, J. Y. Wang, Y. L. Sun, B. Jiang and H. W. Yang, *Chem. Eng. J.*, 2017, **328**, 445-453.
- Z. Q. Zhang, F. Wang, H. H. Nie, T. Zhou, G. D. Zhang, X. F. Wang, M. Cui and Y. Z. Li, *J. Sol-Gel Sci. Technol.*, 2017, **82**, 827-833.
- W. Chen, S. Q. Liang, Y. X. Guo and D. W. Tang, *Energy Convers. Manage.*, 2014, **85**, 13-19.
- W. Chen, S. Q. Liang, Y. X. Guo, X. H. Gui and D. W. Tang, *Fluid Phase Equilib.*, 2013, **360**, 1-6.
- K. S. Lovejoy, C. A. Corley, E. K. Cope, M. C. Valentine, J. G. Leid, G. M. Purdy, J. S. Wilkes, A. T. Koppisch and R. E. Del Sesto, *Cryst. Growth Des.*, 2012, **12**, 5357-5364.
- H. B. Halima, T. Zwingelstein, V. Humblot, B. Lakard and L. Viau, *ACS Appl. Mater. Interfaces*, 2023, **15**, 33382-33396.
- J. Estager, P. Nockemann, K. R. Seddon, M. Swadźba-Kwaśny and S. Tyrrell, *Inorg. Chem.*, 2011, **50**, 5258-5271.
- C. J. Clarke, H. Baaqel, R. P. Matthews, Y. Y. Chen, K. R. J. Lovelock, J. P. Hallett and P. Licence, *Green Chem.*, 2022, **24**, 5800-5812.



18. A. W. Taylor, S. Men, C. J. Clarke and P. Licence, *RSC Adv.*, 2013, **3**, 9436-9445.
19. E. Gousseva, F. K. Towers Tompkins, J. M. Seymour, L. G. Parker, C. J. Clarke, R. G. Palgrave, R. A. Bennett, R. Grau-Crespo and K. R. J. Lovelock, 2024, submitted. View Article Online
DOI: 10.1039/D4FD00029C
20. S. Csihony, H. Mehdi and I. T. Horváth, *Green Chem.*, 2001, **3**, 307-309.
21. J. E. Penner-Hahn, *Coord. Chem. Rev.*, 2005, **249**, 161-177.
22. M. Swadźba-Kwaśny, *RSC Adv.*, 2017, **7**, 51907-51909.
23. T. Waters, X. B. Wang and L. S. Wang, *Coord. Chem. Rev.*, 2007, **251**, 474-491.
24. R. Rowe, K. R. J. Lovelock and P. A. Hunt, *J. Chem. Phys.*, 2021, **155**, 014501.
25. A. Krężel and W. Maret, *Arch. Biochem. Biophys.*, 2016, **611**, 3-19.
26. T. Dudev and C. Lim, *J. Am. Chem. Soc.*, 2000, **122**, 11146-11153.
27. R. R. Roe and Y. P. Pang, *J. Mol. Model.*, 1999, **5**, 134-140.
28. S. I. Hsiu, J. F. Huang, I. W. Sun, C. H. Yuan and J. Shiea, *Electrochim. Acta*, 2002, **47**, 4367-4372.
29. A. P. Abbott, G. Capper, D. L. Davies and R. Rasheed, *Inorg. Chem.*, 2004, **43**, 3447-3452.
30. V. Lecocq, A. Graille, C. C. Santini, A. Baudouin, Y. Chauvin, J. M. Basset, L. Arzel, D. Bouchu and B. Fenet, *New J. Chem.*, 2005, **29**, 700-706.
31. S. Takahashi, L. A. Curtiss, D. Gosztola, N. Koura and M. L. Sabounji, *Inorg. Chem.*, 1995, **34**, 2990-2993.
32. Y. Zou, H. J. Xu, G. Z. Wu, Z. Jiang, S. M. Chen, Y. Y. Huang, W. Huang and X. J. Wei, *J. Phys. Chem. B*, 2009, **113**, 2066-2070.
33. W. H. Wu, Y. X. Lu, Y. T. Liu, H. Y. Li, C. J. Peng, H. L. Liu and W. L. Zhu, *J. Phys. Chem. A*, 2014, **118**, 2508-2518.
34. F. Parveen, T. Patra and S. Upadhyayula, *New J. Chem.*, 2018, **42**, 1423-1430.
35. P. H. Fourcroy, D. Carre and J. Rivet, *Acta Crystallogr. Sect. B: Struct. Sci.*, 1978, **34**, 3160-3162.
36. D. K. Mishra, G. Gopakumar, G. Pugazhenthii, C. Rao, S. Nagarajan and T. Banerjee, *J. Phys. Chem. A*, 2021, **125**, 9680-9690.
37. A. P. Abbott, J. C. Barron, G. Frisch, S. Gurman, K. S. Ryder and A. F. Silva, *Phys. Chem. Chem. Phys.*, 2011, **13**, 10224-10231.
38. K. R. J. Lovelock, I. J. Villar-Garcia, F. Maier, H. P. Steinrück and P. Licence, *Chem. Rev.*, 2010, **110**, 5158-5190.
39. K. R. J. Lovelock and P. Licence, in *Ionic Liquids UNCOILED: Critical Expert Overviews*, eds. K. R. Seddon and N. V. Plechkova, Wiley, Oxford, 2012, ch. 8, pp. 251-282.
40. H. P. Steinrück, *Phys. Chem. Chem. Phys.*, 2012, **14**, 5010-5029.
41. M. L. Baker, M. W. Mara, J. J. Yan, K. O. Hodgson, B. Hedman and E. I. Solomon, *Coord. Chem. Rev.*, 2017, **345**, 182-208.
42. F. Maier, T. Cremer, C. Kolbeck, K. R. J. Lovelock, N. Paape, P. S. Schulz, P. Wasserscheid and H. P. Steinrück, *Phys. Chem. Chem. Phys.*, 2010, **12**, 1905-1915.
43. P. A. Brühwiler, O. Karis and N. Mårtensson, *Rev. Mod. Phys.*, 2002, **74**, 703-740.
44. J. L. Campbell and T. Papp, *At. Data Nucl. Data Tables*, 2001, **77**, 1-56.
45. D. Briggs and J. T. Grant, eds., *Surface Analysis by Auger and X-ray Photoelectron Spectroscopy*, IM Publications, Manchester, 2003.
46. J. C. Vickerman and I. S. Gilmore, eds., *Surface Analysis: The Principal Techniques*, John Wiley & Sons, Chichester, 2009.
47. P. S. Bagus, E. S. Ilton and C. J. Nelin, *Surf. Sci. Rep.*, 2013, **68**, 273-304.
48. E. Gousseva, S. D. Midgley, J. M. Seymour, R. Seidel, R. Grau-Crespo and K. R. J. Lovelock, *J. Phys. Chem. B*, 2022, **126**, 10500-10509.
49. J. Liu, H. Zhang, Y. M. Li and Z. Liu, *J. Phys. Chem. B*, 2018, **122**, 10600-10606.
50. R. A. Walton, *Coord. Chem. Rev.*, 1976, **21**, 63-91.
51. J. M. Seymour, E. Gousseva, A. I. Large, C. J. Clarke, P. Licence, R. M. Fogarty, D. A. Duncan, P. Ferrer, F. Venturini, R. A. Bennett, R. G. Palgrave and K. R. J. Lovelock, *Phys. Chem. Chem. Phys.*, 2021, **23**, 20957-20973.
52. S. Hayama, G. Duller, J. P. Sutter, M. Amboage, R. Boada, A. Freeman, L. Keenan, B. Nutter, L. Cahill, P. Leicester, B. Kemp, N. Rubies and S. Diaz-Moreno, *J. Synchrotron Radiat.*, 2018, **25**, 1556-1564.
53. S. Hayama, R. Boada, J. Chaboy, A. Birt, G. Duller, L. Cahill, A. Freeman, M. Amboage, L. Keenan and S. Diaz-Moreno, *J. Phys.: Condens. Matter*, 2021, **33**, 11.
54. P. Glatzel, M. Sikora, G. Smolentsev and M. Fernandez-Garcia, *Catal. Today*, 2009, **145**, 294-299.
55. M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. J. A. Montgomery, J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Journal*, 2016.
56. J. D. Chai and M. Head-Gordon, *J. Chem. Phys.*, 2008, **128**, 084106.
57. J. D. Chai and M. Head-Gordon, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6615-6620.
58. F. Weigend, *Phys. Chem. Chem. Phys.*, 2006, **8**, 1057-1065.
59. F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297-3305.
60. V. S. Bernales, A. V. Marenich, R. Contreras, C. J. Cramer and D. G. Truhlar, *J. Phys. Chem. B*, 2012, **116**, 9122-9129.
61. F. Neese, *Wiley Interdiscip. Rev.-Comput. Mol. Sci.*, 2012, **2**, 73-78.
62. A. Najibi and L. Goerigk, *J. Chem. Theory Comput.*, 2018, **14**, 5725-5738.
63. E. van Lenthe, J. G. Snijders and E. J. Baerends, *J. Chem. Phys.*, 1996, **105**, 6505-6516.
64. N. Fairley, V. Fernandez, M. Richard-Plouet, C. Guillot-Deudon, J. Walton, E. Smith, D. Flahaut, M. Greiner, M. Biesinger, S. Tougaard, D. Morgan and J. Baltrusaitis, *Appl. Surf. Sci. Adv.*, 2021, **5**, 100112.
65. C. Kolbeck, M. Killian, F. Maier, N. Paape, P. Wasserscheid and H. P. Steinrück, *Langmuir*, 2008, **24**, 9500-9507.
66. T. Cremer, C. Kolbeck, K. R. J. Lovelock, N. Paape, R. Wölfel, P. S. Schulz, P. Wasserscheid, H. Weber, J. Thar, B. Kirchner, F. Maier and H. P. Steinrück, *Chem.-Eur. J.*, 2010, **16**, 9018-9033.
67. I. J. Villar-Garcia, E. F. Smith, A. W. Taylor, F. L. Qiu, K. R. J. Lovelock, R. G. Jones and P. Licence, *Phys. Chem. Chem. Phys.*, 2011, **13**, 2797-2808.
68. L. G. Parker, F. K. Towers Tompkins, J. M. Seymour, N. O. Alblewi, E. Gousseva, M. R. Daw, S. Hayama, R. P. Matthews, A. A. E. Fouda, J. D. Elliott, C. D. Smith and K. R. J. Lovelock, 2024, in preparation.



69. H. A. Øye, M. Jagtoyen, T. Oksefjell and J. S. Wilkes, *3rd International Symp on Molten Salt Chemistry and Technology (Ms3 Symp)*, 1991, **73**, 183-189.
70. O. G. Parchment, M. A. Vincent and I. H. Hillier, *J. Phys. Chem.*, 1996, **100**, 9689-9693. View Article Online
DOI: 10.1039/D4FD00029C
71. C. C. Pye, C. R. Corbeil and W. W. Rudolph, *Phys. Chem. Chem. Phys.*, 2006, **8**, 5428-5436.
72. C. C. Pye, S. M. Black and W. W. Rudolph, *J. Solution Chem.*, 2011, **40**, 1932-1954.
73. L. G. Parker, F. K. Towers Tompkins, J. M. Seymour, E. Gousseva, N. O. Alblewi, C. A. Borrill, C. J. Clarke, S. Hayama, R. Seidel, D. Céolin, R. H. Temperton, R. P. Matthews, C. D. Smith and K. R. J. Lovelock, 2024, in preparation.
74. S. B. Pillai, R. J. Wilcox, B. G. Hillis, B. P. Losey and J. D. Martin, *J. Phys. Chem. B*, 2022, **126**, 2265-2278.
75. T. M. Laher and C. L. Hussey, *Inorg. Chem.*, 1983, **22**, 3247-3251.

