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Geometries, interaction energies and bonding in $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ and $[\text{PoCl}_n]^{4-n}$ complexes†

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Polonium ($Z = 84$) is one of the rarest elements on Earth. More than a century after its discovery, its chemistry remains poorly known and even basic questions have not yet been satisfactorily addressed. In this work, we perform a systematic study of the geometries, interactions energies and bonding in basic polonium(IV) species, namely the hydrated $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ and chlorinated $[\text{PoCl}_n]^{4-n}$ complexes by means of gas-phase electronic structure calculations. We show that while up to nine water molecules can fit in the first coordination sphere of the polonium(IV) ion, its coordination sphere can already be filled with eight chloride ligands. Capitalising on previous theoretical studies, a focused methodological study based on interaction energies and bond distances allows us to validate the MP2/def2-TZVP level of theory for future ground-state studies. After discussing the similarities and differences between complexes with the same number of ligands, we perform topological analyses of the MP2 electron densities in the quantum theory of atoms in molecules (QTAIM) fashion. While the water complexes display typical signatures of closed-shell interactions, we reveal large Po–Cl delocalisation indices, especially in the hypothetical $[\text{PoCl}]^{3+}$ complex. This “enhanced” covalency opens the way for a significant spin–orbit coupling (SOC) effect on the corresponding bond distance, which has been studied using two independent approaches (*i.e.* one *a priori* and one *a posteriori*). We finally conclude by stressing that while the SOC may not affect much the geometries of high-coordinated polonium(IV) complexes, it should definitely not be neglected in the case of low-coordinated ones.

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1 Introduction

Since the discovery of polonium in 1898 by Marie and Pierre Curie, not a lot of experiments have been carried out to unravel its intriguing chemical properties. While most of polonium chemistry remains unknown, its extreme toxicity, being either chemical or radiological, is notorious.^{1,2} Though its radioactivity may be used to detect it, it is still complicated to perform analyses to determine polonium traces in, for example, sea water,³ tobacco,^{4,5} plants,⁶ soils⁷ and other samples. Meanwhile, former usages of polonium as a heat source in the space machine industry, as an α -emission source, or as static eliminators have now become obsolete because of alternatives based on

materials that are safer and easier to obtain. In the case of accidental or criminal ingestion of polonium, no specific antidote is known, and one can administrate British anti-Lewisite (BAL), the antidote for heavy metals and in particular arsenic,⁸ unfortunately without any guarantee of success.

Polonium has more than 40 known isotopes. While only seven of them are naturally occurring,⁹ ^{210}Po is the usual isotope for performing experiments, even if other ones may be scarcely used, *e.g.* ^{208}Po for atomic spectroscopy.¹⁰ The four oxidation states of polonium usually considered are: $-II$, $+II$, $+IV$ and $+VI$. In solution, polonium(IV) is meant to be the most stable one,¹¹ so it is clearly an oxidation state whose chemistry deserves to be studied. Nonetheless, evidence for polonides (Na_2Po , ZnPo , *etc.*),¹² oxides (PoO_2 , PoO_3),¹³ halides and dihalides (PoCl_2 , PoCl_4 , PoBr_4 , and PoI_4)^{14–16} and complexes with organic ligands such as the acetylacetonate anion (acac^-), the ethylenediaminetetraacetic acid trianion (EDTA^{3-}), and the oxalate dianion ($\text{C}_2\text{O}_4^{2-}$), *etc.* has been reported in the literature.¹⁷

Computational chemistry is, of course, free of any direct toxicological impact. Its increased accuracy and applicability observed in the last few decades may open the way for future predictions aiming at designing new successful experiments with polonium. However, prior to this, methodological studies have to be performed in order to ensure good results for good

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reasons (or at least identified error cancellations). In fact, only a limited number of electronic structure theory studies have been dedicated to polonium, among which a few emblematic ones will be discussed hereafter. Note that the electronic structure of the free Po^{4+} ion is $[\text{Xe}]4f^{14}5d^{10}6s^2$, which has two main consequences for the calculations, (i) single-configurational and single-reference approaches may correctly describe the ground-state wave functions of polonium(IV) species and (ii) relativistic effects must be accounted for explicitly or implicitly (*e.g.* scalar relativistic effects lead to a strong stabilisation of the 6s orbital¹⁸).

From 2008 to 2010, Ayala *et al.* published three computational works related to polonium(IV) hydration and to the formation of hydroxo complexes in solution.^{19–21} In 2015, Borschevsky *et al.* calculated very accurately with a fully relativistic coupled-cluster method the ionisation potentials and electron affinity of the free Po atom.²² Note that their first ionization potential value was in good accord with the early value of Charles,¹⁰ and also proved to be in good accord with the yet to come new experimental values derived from two independent experiments.^{23,24} In 2019, Stoianov *et al.* revisited from theory the UV-vis absorption spectra of polonium chloride complexes.²⁵ The same year, chalcogen bonding with polonium was examined by Zierkiewicz *et al.*²⁶ In fact, performing relativistic and correlated calculations is always quite challenging, and validating approximate schemes from scratch may require tedious methodological studies due to the lack of experimental reference data (one should then validate more approximate schemes by comparison with more accurate ones, *i.e.* based on very demanding calculations, see for instance the work of Rota *et al.* on the polonium hypothetical dimer²⁷).

As mentioned before, the +IV oxidation state of polonium dominates in solution. Following the pioneering works of Ayala *et al.*¹⁹ and Stoianov *et al.*,²⁵ we will focus on two families of Po(IV)-complexes, namely the hydrated $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ and the chlorinated $[\text{PoCl}_n]^{4-n}$ complexes. The first series is of fundamental interest since it relates to the general context of the hydration of heavy-element cations in solution.²⁸ The second series is of practical interest since polonium chloride complexes can form in acidic chloride solutions.^{12,28} We will focus on ground-state properties (geometries, interaction energies and bonding descriptors) and assess the dependence of the theoretical results with respect to specific degrees of freedom such as the choice of the correlation method, of the atomic basis sets or for instance of correcting for the basis set superposition error (BSSE) when computing interaction energies, which was not fully done in those earlier works.^{19,25} To limit the discussion to pure electronic-structure theory aspects, only gas-phase results will be considered (future work may deal with solvation). Similarities and differences between the hydrated and chlorinated complexes will be discussed, in particular concerning the first coordination sphere of the polonium(IV) cation.

2 Computational details

Our most extended methodological investigation was done on the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ complexes (where in fact $n = 1–9$), even though

some tests were repeated on the $[\text{PoCl}_n]^{4-n}$ ones ($n = 1–8$, see the Results and discussion section and the ESI†). MP2^{29–33} and B3LYP^{34,35} calculations were performed to determine the molecular geometries and the corresponding total interaction energies of all the complexes. Because of the close agreement of the MP2 and DFT (B3LYP) results and for the sake of simplicity, we will only report MP2 results in the remainder of the text. For the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ ($n = 7–9$) complexes, initial structures were built from previous thorium structures.³⁶ Note that additional reference calculations with the coupled cluster method with single, double and perturbative triple excitations,^{37,38} *i.e.* CCSD(T), were also performed on specific complexes, namely the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ with $n = 1–3$. All the MP2, B3LYP and CCSD(T) calculations were carried out with Gaussian 16³⁹ software. Note that every molecular geometry is optimised unless mentioned otherwise.

The Dunning⁴⁰ aug-cc-pVDZ, aug-cc-pVTZ, and aug-cc-pVQZ, and redefined Ahlrichs⁴¹ def2-TZVP and def2-QZVP basis sets together with the relativistic 60-electron small core ECP60MDF pseudopotential⁴⁰ for Po and its associated aug-cc-pVNZ-PP or def2-NZVP-PP basis sets were used ($N = \text{D, T, Q}$). All the basis sets and pseudopotential parameters were taken from the Basis Set Exchange software.⁴²

By default, in post-Hartree-Fock calculations, Gaussian 16 correlates all outer-core (5s, 5p, and 5d) orbitals and the occupied 6s valence orbital of Po(IV), while freezing the 5s and 5p orbitals is wiser to ensure a significant energy gap between the frozen and the correlated orbitals in a molecular system, as illustrated by Fig. 1 related to the PoCl_4 molecule, where the Po 5s molecular orbital (MO 21) is energetically close to the frozen combinations of the chlorine 2s (MOs 9–12) and 2p orbitals (MOs 13–20). To generally avoid artefacts in the correlated calculations (when discussing trends, one should only compare results for which the same shells are frozen), the default frozen core choice was overwritten with the window

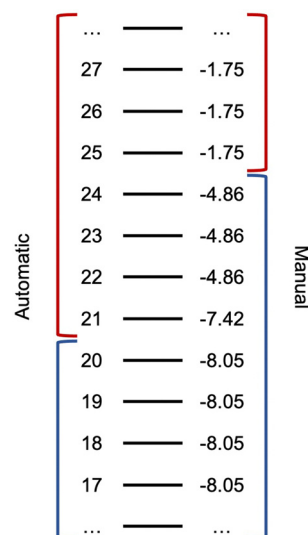


Fig. 1 Hartree-Fock/def2-TZVP orbital energies (in a.u.) of the PoCl_4 complex. The frozen (blue) and unfrozen (red) orbitals are displayed, by default (left) or with our recommended scheme (right).



keyword of Gaussian, with which the user may define the first unfrozen orbital, *i.e.* the number of frozen cores (NFC) plus one. In fact, in the current study, it was manually set to freeze 4 orbitals for Po (5s and 5p pseudo-orbitals), 1 for each water (the 1s orbital of the O atom), and 5 for Cl (the 1s, 2s and 2p shells). Moreover, though it is a bit technical, we would like to warn the reader that attention should be paid to compute the frequencies in the second step to ensure that Gaussian is correctly applied to the desired freezing scheme.

The interaction energies were calculated using the following formula:

$$E_{\text{int}} = E_{\text{complex}} - E_{\text{Po}^{4+}} - nE_{\text{ligand}} + \Delta E_{\text{BSSE}} \quad (1)$$

The basis set superposition error correction (ΔE_{BSSE}) was calculated by means of the counterpoise correction at the (fixed) geometry of the complex:

$$\Delta E_{\text{BSSE}} = E_{\text{Po}^{4+}} - E_{\text{Po}^{4+}}^{\text{ghosts}} - E_{(\text{ligands})_n} + E_{(\text{ligands})_n}^{\text{ghost}} \quad (2)$$

where the ghost(s) superscript refers to calculations performed with ghost ligand atoms ($E_{\text{Po}^{4+}}^{\text{ghosts}}$) or with a ghost Po atom ($E_{(\text{ligands})_n}^{\text{ghost}}$), meaning that the basis functions of the corresponding atoms are applied but their electrons are not (neither their nuclear charges). Note that unlike eqn (1), which dissociates the complex in $\text{Po}^{4+} + n$ ligands, eqn (2) considers clusters of n interacting ligands, denoted $(\text{ligands})_n$.

To describe the nature of the chemical bonds in the systems under study, quantum theory of atoms in molecules (QTAIM)⁴³ analyses were done with the AIMALL⁴⁴ software using the MP2/def2-TZVP electron densities. Note that more details concerning the meaning of the derived descriptors will be directly given in the Results and discussion section.

To assess the role of the spin-orbit coupling (SOC) on molecular geometries, two types of calculations were performed. First, zeroth-order regular approximation (ZORA)⁴⁵ relativistic Hartree-Fock calculations were performed using the AMS driver of the Amsterdam Modeling Suite⁴⁶ software and Slater-type TZ2P⁴⁷ basis sets, without freezing core orbitals. In this case, the SOC is included *a priori*, but the interplay between SOC and electron correlation is completely missing. Alternatively, we have performed two-step SOC calculations with ORCA⁴⁸ (v. 5.0.0), for which the SOC is introduced *a posteriori*, as a perturbation of the correlated spin-orbit free (SOF) level, obtained with the Douglas-Kroll-Hess Hamiltonian.^{49–51} In this case, specific scalar relativistic recontracted (DKH-def2-TZVP) or originally generated (SARC-DKH-TZVP) all-electron basis sets^{52,53} were used for all the atoms. There, the role of the SOC may this time be underestimated, and only a part of the interplay between electron correlation and the SOC is by construction treated (we have only used the contracted spin-orbit configuration interaction, SOCI, scheme). More details and explanation will be directly given in the Results and discussion section. In short, these two complementary approaches do not suffer from the same pitfalls, and if a physical

effect is observed with both methodologies, one may thus be quite confident in its actual occurrence.

3 Results and discussion

3.1 Electronic structure theory calculations

As mentioned earlier, we will start the discussion with the interactions energies that were obtained for the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ and $[\text{PoCl}_n]^{4-n}$ complexes, with special emphasis on the water complexes, previously studied by Ayala *et al.*¹⁹ We performed a partial benchmark to find the best accuracy/cost methodology for ground-state studies of polonium(IV) complexes. We recall here some of their important conclusions, confirmed by our independent calculations:

- Up to nine water molecules can fit in the polonium(IV) coordination sphere, meaning that $n = 1-9$ for the hydrated $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ complexes under study.
- The interaction energies evolve monotonously with n .
- MP2 and to a lesser extend B3LYP calculations provide interaction energies in good agreement with the CCSD(T) ones when the aug-cc-pVDZ basis set are used.

We have thus retained the MP2 method, and focused on two main degrees of freedom: the nature and size of the basis set and the use of a BSSE correction (see eqn (1) and (2)). Furthermore, since we have proposed to change the number of polonium frozen orbitals (see the Computational details section), we have also reported new CCSD(T) and MP2 calculations with the aug-cc-pVDZ basis set, as well as original ones with triple zeta and quadruple zeta aug-cc or def2 basis sets (see Table 1, as well as Tables S1 and S2, ESI† for an investigation of the impact of the BSSE correction).

One can quickly see that results for both the MP2 and CCSD(T) methods, in combination with the aug-cc-pVDZ basis set, are in fair agreement, though the accord is less impressive than in the paper of Ayala *et al.*¹⁹ Since the BSSE corrections are similar with both methods (see Tables S1 and S2, ESI†), the observed difference essentially comes from the number of frozen orbitals. However, it is known that wave-function theory calculations, especially correlated ones of the coupled-cluster type, slowly converge with the basis-set size. Because of the

Table 1 Interaction energies with BSSE correction (kcal mol^{-1}) obtained at various levels of theory for polonium(IV) complexes with water and chlorides, computed according to eqn (1) and (2)

n (ligands)	H_2O				Cl^-	
	CCSD(T)/ aug-cc- pVDZ	MP2/aug- cc-pVDZ	MP2/aug- cc-pVTZ	MP2/def2- TZVP	MP2/def2- QZVP	MP2/ def2- TZVP
1	−236.2	−249.2	−257.9	−262.0	−263.8	−883.7
2	−409.5	−439.0	−454.7	−461.4	−464.9	−1415.0
3	−541.1	−586.4	−607.7	−616.1	−621.4	−1749.2
4		−694.9	−718.8	−728.9		−1911.4
5		−791.8	−817.6	−829.4		−1974.3
6		−875.8	−903.1	−918.2		−1947.8
7		−941.2	−971.1	−987.8		−1861.4
8		−1004.4	−1035.9	−1054.0		−1650.3
9		−1052.7	−1085.5	−1104.4		



computational cost, we have not systematically performed CCSD(T) calculations with larger basis sets and have essentially investigated the effect of the basis-set nature and size only at the MP2 level. However, the interested reader may find a couple of CCSD(T) optimised structures obtained with the aug-cc-pVTZ and def2-TZVP basis sets in Table S2 (ESI†). Interestingly, the accord between MP2 and CCSD(T) improves with basis set enlargement, independent from its nature (*i.e.* aug-cc *vs.* def2). Indeed, the difference between the two methods is divided by two with respect to the basis-set size. Anyway, due to the single reference nature of the ground state of polonium(IV) complexes, we indeed expect to obtain accurate values with MP2 provided that convergence of the results with respect to the basis-set size is reached. This is why we chose to pursue analysis of the results dependence with respect to the basis set with solely the MP2 method.

Nowadays, computational facilities allow us to proceed with larger basis sets than fifteen years ago. For instance, we have reported calculations with the aug-cc-pVTZ, def2-TZVP and def2-QZVP basis sets (see Table 1). By comparing the aug-cc-pVDZ and aug-cc-pVTZ results, it is clear that the MP2 interaction energies are not yet converged with the aug-cc-pVDZ basis set, which somehow justifies *a posteriori* the present benchmark. Thus, at least a triple-zeta basis set should be used. As an alternative to the aug-cc basis sets, we have employed the def2-TZVP and def2-QZVP ones (see Table 1). Notably, the def2-TZVP basis set shows very similar results to the ones obtained with the aug-cc-pVTZ basis set, despite being built with fewer functions. The largest difference between the corresponding sets of results is below 20 kcal mol⁻¹ for the $n = 9$ complex, which translates into ~ 2 kcal mol⁻¹ per ligand. Since the calculations with the aug-cc-pVTZ basis set take roughly 5 to 20 times more time than the def2-TZVP one, it is clear that the best accuracy/cost ratio is expected for the def2 basis sets, in particular the def2-TZVP one.

As an additional step of analysis, we compared the results obtained with the def2-TZVP and def2-QZVP basis sets. As can be seen in the ESI† (Tables S1 and S2), interaction energies differ by less than 2 kcal mol⁻¹ per ligand when the BSSE correction is included (the correction effectively reducing this difference). Considering the higher computational cost of calculations with the def2-QZVP basis set, we conclude that the MP2/def2-TZVP level of theory reaches the best accuracy to cost ratio out of all the tested methods for computing interaction energies between the Po⁴⁺ ion and the coordinated water ligands. Furthermore, we do not expect a notable improvement of the interactions energies by performing a complete basis-set extrapolation^{54–56} based on the def2-TZVP and def2-QZVP calculations.

We have repeated similar calculations on the chlorinated complexes. First, we have found that eight chloride ligands are sufficient to saturate the coordination sphere of polonium(IV). Second, we observe a change in the evolution of the interaction energies (see Table 1), which is *a priori* more visible in Fig. 2. While in the case of water, the evolution is monotonous with respect to ligand addition, the curve for chloride ligands

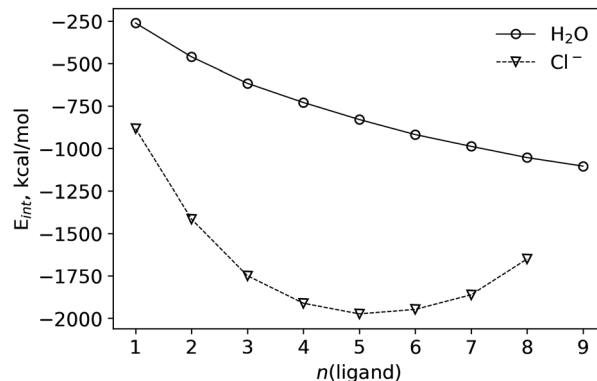


Fig. 2 MP2/def2-TZVP interaction energies (kcal mol⁻¹), E_{int} , as functions of the number of ligands (n) for polonium(IV) complexes with water and chlorides.

displays a minimum for $n = 5$ (note that all the structures are stable though). We connect this phenomenon with the negative charges of chloride ligands: for each ligand addition, we earn a coordination bond but we also enhance the inter-ligand repulsion in a much stronger way than with neutral ligands. We have also repeated a similar benchmark as what was done for the stepwise water addition (see Tables S3 and S4, ESI†), leading to the exact same conclusion: the MP2/def2-TZVP level of theory is a good compromise to compute accurate interaction energies.

As a part of our benchmark study, we have been looking at molecular geometries, with emphasis on the mean Po–O and Po–Cl bond lengths. Prior to fully digging into this, we start with an overall description of the structures. Perspective views are given in Fig. 3, together with the symmetry point groups (SPGs). For the polonium(IV) complexes with water, we obtain similar structures as the ones reported by Ayala *et al.*,¹⁹ if we exclude the $[\text{Po}(\text{H}_2\text{O})_7]^{3+}$ complex from the discussion (a different SPG may be found depending on the applied level of theory). Concerning this specific complex, note that we report a C_2 SPG, obtained at the MP2/def2-TZVP level of theory.

When comparing the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ and $[\text{PoCl}_n]^{4-n}$ structures two-by-two for $n = 1$ –8, several situations occur:

- For $n = 2$, the same SPG is observed.
- For $n = 1, 3, 6$ and 8 , a direct subgroup of the SPG of the $[\text{PoCl}_n]^{4-n}$ complex is found for the corresponding $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ complex.
- For $n = 4, 5$ and 7 , the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ complexes display, if any, much less symmetry elements than the corresponding $[\text{PoCl}_n]^{4-n}$ ones. Those structures are thus considered as qualitatively different.

Though it is obvious that the presence of the hydrogen atoms of the water ligands may lower the symmetry, it is actually interesting to note that in some of the cases ($n = 4, 5$ and 7) quite different structures were obtained.

Concerning more specifically the PoCl_4 complex, Stoianov *et al.* reported in their article²⁵ a C_{2v} SPG for the gas-phase structure, with one set of bonds being slightly longer than the other. In fact, we found with our level of theory that the gas phase C_{2v} structure is only 1.6 kcal mol⁻¹ higher in energy than



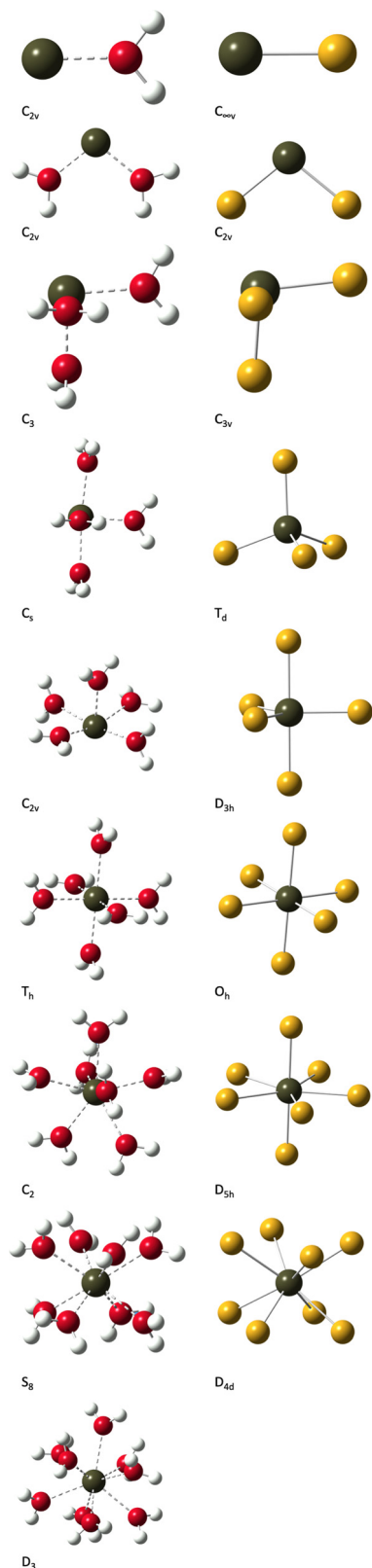


Fig. 3 Ball-and-stick representations and symmetry point groups of the studied polonium(IV) complexes with water molecules and chlorides. Colour code: polonium (dark green), chlorine (yellow), oxygen (red) and hydrogen (white).

Table 2 Mean MP2 Po–O and Po–Cl bond lengths (Å), obtained for polonium(IV) complexes with water molecules and chlorides, respectively

<i>n</i> (ligands)	H ₂ O			Cl [−]
	aug-cc-pVDZ	aug-cc-pVTZ	def2-TZVP	def2-TZVP
1	2.08	2.04	2.03	2.25
2	2.11	2.08	2.07	2.28
3	2.15	2.11	2.11	2.34
4	2.2	2.17	2.17	2.47
5	2.25	2.22	2.22	2.54
6	2.30	2.27	2.27	2.60
7	2.34	2.31	2.31	2.72
8	2.37	2.34	2.34	2.83
9	2.41	2.38	2.39	

the T_d one, meaning that both are technically possible. We have thus performed several test calculations with various levels of theory, and concluded that though the potential energy hyper-surface may be quite flat, the T_d structure must be the correct gas-phase one (we recall here that a structure similar to the one of the corresponding water complex is expected when a solvation model is applied²⁵). Finally, we would like to mention that the structure of the $[\text{Po}(\text{H}_2\text{O})_9]^{4+}$ complex, which has no match in the chlorinated complex series, is highly symmetric (D_3), with 6 and 3 by symmetry-related bonds, respectively.

The dependence of the mean Po–O and Po–Cl bond lengths with respect to the basis-set sizes may be seen from Table 2, Table S5 and S6 (ESI[†]), and is also displayed in Fig. 4 for the polonium(IV) complexes with water. In line with the interaction energies, we note significant shortening of the bonds by going from the aug-cc-pVDZ basis set to the aug-cc-pVTZ one, a good agreement between the aug-cc-pVTZ and def2-TZVP results, and little change by going from def2-TZVP to def2-QZVP (see Table S5, ESI[†]), meaning that the def2-TZVP basis set is again a good accuracy/cost ratio performer.

For the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ ($n = 1-3$) complexes, we obtain similar MP2/aug-cc-pVDZ bond distances to the ones of Ayala *et al.*,¹⁹ meaning that our proposed change in the NFCs has virtually no effect on the molecular geometries. Consequently, the good agreement of the MP2 and CCSD(T) structures that they

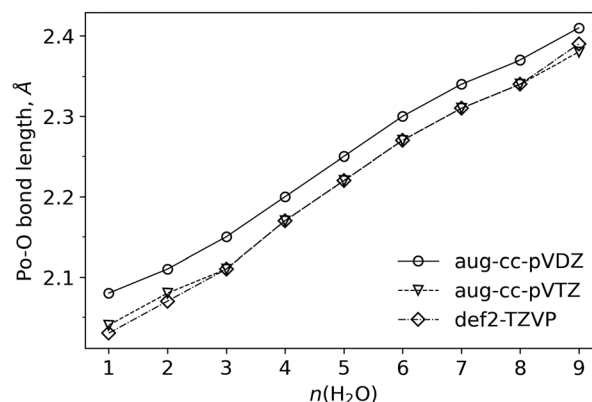


Fig. 4 Mean MP2 Po–O bond lengths (Å), obtained for polonium(IV) complexes with water for various basis sets, as functions of the number of water molecules (*n*).



observed is preserved in the present study. Concerning the $[\text{PoCl}_n]^{4-n}$ ($n = 4-6$) complexes, the mean B3LYP/aug-cc-pVTZ bond distances reported by Stoianov *et al.*²⁵ (2.52, 2.59 and 2.67, respectively), are significantly longer than the MP2/def2-TZVP ones (by ~ 0.05 Å), which is in line with the ~ 0.03 Å observed by Ayala *et al.*¹⁹ for the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ ($n = 1-9$) series. Note that the application of a dispersion correction⁵⁷ slightly diminishes those bond distances (by ~ 0.01 Å in the $[\text{PoCl}_n]^{4-n}$, $n = 4-6$, complexes), which thus only explains part of the difference between B3LYP and MP2 (see Table S6, ESI†). Since the bond lengths of the $[\text{Po}(\text{H}_2\text{O})_n]^{4+}$ ($n = 1-3$) complexes tend to be overestimated by ~ 0.02 Å,¹⁹ we conclude that our new MP2/def2-TZVP structures for the $[\text{PoCl}_n]^{4-n}$ ($n = 4-6$) complexes are actually more accurate than the previously reported ones.²⁵

3.2 QTAIM analyses

To better characterise the nature of the bonds in the studied polonium complexes, QTAIM⁴³ analyses were performed. Several indicators may be defined, in particular at the bond critical points (BCPs). Such indicators may be used to classify the nature of the bonds, as proposed by Nakanishi and Hayashi⁵⁸ and validated by Pilmé *et al.*⁵⁹ for astatine species (a neighbour of polonium in the periodic table). We report for convenience the bond lengths, and more specifically the electron density values at the BCPs (ρ), the Laplacian of the density at the BCP ($\nabla^2\rho$), the kinetic energy density (G), the potential energy density (V), the ratio between the absolute value of the potential energy density to the kinetic energy one ($|V|/G$), the delocalisation indices (δ), and the Po and O or Cl atomic charges (q) in Tables 3 and 4.

With the value of the electron density at the BCP of interest, one can determine the nature of the bond, in particular, determine if the interaction is of the shared-shell ($\rho \geq 0.2$ a.u.) or closed-shell type. This classification may also be done based on the following two relationships for the Laplacian ($\nabla^2\rho(r)$) and the total electron energy densities at BCPs ($H(r)$):⁵⁸

$$\frac{1}{4}\nabla^2\rho(r) = 2G(r) + V(r) \quad (3)$$

$$H(r) = G(r) + V(r) \quad (4)$$

where $G(r)$ is positive and $V(r)$ is negative. Actually, if the Laplacian of the density at the BCP is positive, as in all the studied complexes, then the kinetic energy density dominates, meaning that we are dealing with a closed-shell interaction.⁵⁸

Another criterion that can help us with the description of chemical bonds is the ratio between the absolute value of the potential energy density to the kinetic energy one. If this ratio is smaller than 1, we expect a pure closed-shell interaction (thus a pure ionic bond or a typical non-covalent interaction). If this ratio is larger than 1, which is the case in all the studied complexes, it means that the potential energy is large (electrons are thus stabilized at the BCP⁵⁸), and we face a regular closed-shell interaction.

For both types of complexes we can see the same trend: the larger the complex, the longer the bond, the smaller the values of electron density at the BCP, and so do the values of the Lagrangian, potential and kinetic energy densities. This is somehow logical since the addition of a ligand will lead to smaller charge transfers from each ligand to the polonium(IV) ion, *i.e.* to reduced “covalency”, and since inter-ligand repulsion is also enhanced (which obviously is much more pronounced for chloride ligands, as already mentioned).

The delocalisation index (δ)^{60,61} is also another important criterion, meant to be related to the bond multiplicity. A plot of the comparison between the mean δ (Po–O) and δ (Po–Cl) values is depicted in Fig. 5, allowing a quicker view at the general trends than by inspecting Tables 3 and 4. First, we can see that almost all the values are lower or equal to 1, which represents an ideal single bond (the two atoms exactly sharing one electron pair). In fact, half of the complexes presented here have δ 's that are smaller than 0.5, meaning that the corresponding bonds are far from being covalent. Since a δ value around 1 may be indicative of several bonding patterns (*i.e.* covalent, charge shift⁶² or even iono-covalent), these are not incompatible with our previous conclusion (regular closed-shell interactions). In fact, the more striking value is the one

Table 3 MP2/def2-TZVP Po–O bond distances (Å) and QTAIM bonding descriptors for polonium(IV) complexes with water molecules

n (H_2O)	r (Po–O), Å	ρ , a.u.	$\nabla^2\rho$	G , a.u.	V , a.u.	$ V /G$	δ (Po–O)	q (Po), a.u.	q (O), a.u.	q (H), a.u.
1	2.03	0.20	0.42	0.15	−0.20	1.30	1.1	3.36	−1.04	0.84
2	2.07	0.12	0.36	0.13	−0.17	1.31	0.8	3.03	−1.13	0.80
3	2.11	0.11	0.31	0.11	−0.15	1.31	0.6	2.91	−1.20	0.78
4	2.13	0.10	0.29	0.10	−0.14	1.31	0.5	2.87	−1.23	0.77
	2.22 (x2)	0.08	0.24	0.08	−0.11	1.28	0.6		−1.25	
	2.12	0.10	0.30	0.11	−0.14	1.31	0.6		−1.21	
5	2.13	0.10	0.30	0.10	−0.13	1.29	0.4	2.85	−1.23	0.75
	2.24 (x2)	0.08	0.22	0.08	−0.10	1.27	0.5		−1.26	0.74
	2.24 (x2)	0.08	0.22	0.08	−0.10	1.27	0.5		−1.26	0.74
6	2.27	0.07	0.21	0.07	−0.09	1.26	0.4	2.84	−1.26	0.73
7	2.32 (x2)	0.07	0.18	0.06	−0.07	1.23	0.3	2.88	−1.27	0.71
	2.34 (x2)	0.06	0.18	0.06	−0.07	1.23	0.3		−1.27	0.72
	2.29	0.07	0.20	0.06	−0.08	1.24	0.3		−1.26	
	2.28 (x2)	0.07	0.20	0.07	−0.08	1.24	0.4		−1.26	
8	2.34	0.06	0.18	0.06	−0.07	1.21	0.3	2.9	−1.27	0.70
9	2.43 (x3)	0.05	0.15	0.04	−0.05	1.17	0.2	2.92	−1.26	0.69



Table 4 MP2/def2-TZVP Po–Cl bond distances (Å) and QTAIM bonding descriptors for polonium(IV) complexes with chlorides

n (Cl [−])	r (Po–Cl), Å	ρ , a.u.	$\nabla^2\rho$	G , a.u.	V , a.u.	$ V /G$	δ (Po–Cl)	q (Po), a.u.	q (Cl), a.u.
1	2.25	0.12	0.26	0.12	−0.17	1.46	1.8	2.52	0.48
2	2.28	0.12	0.18	0.10	−0.15	1.53	1.2	2.06	−0.03
3	2.34	0.11	0.13	0.08	−0.12	1.57	0.9	1.94	−0.31
4	2.47	0.08	0.13	0.06	−0.09	1.46	0.7	1.98	−0.50
5	2.56 (x2)	0.07	0.11	0.05	−0.07	1.43	0.6	2.06	−0.64
	2.53 (x3)	0.07	0.11	0.05	−0.07	1.44	0.6		−0.59
6	2.60	0.06	0.10	0.04	−0.06	1.41	0.5	2.13	−0.69
7	2.75 (x5)	0.05	0.07	0.03	−0.04	1.34	0.4	2.13	−0.74
	2.63 (x2)	0.06	0.09	0.04	−0.05	1.39	0.5		−0.71
8	2.83	0.04	0.07	0.02	−0.03	1.28	0.4	2.12	−0.77

observed for the [PoCl]³⁺ complex: 1.8. This value somehow resembles what is expected for a double bond, meaning that this bond must bear a strong covalent character.

In line with the delocalisation index, the QTAIM atomic q (Cl) charge in [PoCl]³⁺ is positive (strong Cl[−] to Po⁴⁺ charge transfer). Knowing that atomic charges are strongly dependent on the underlying charge model, we have also computed the Mulliken⁶³ and Hirshfeld⁶⁴ atomic charges (see Table S7, ESI[†]). Since all the applied charge models lead to a positive q (Cl) in [PoCl]³⁺, we conclude for a strong charge transfer from Cl[−] to Po⁴⁺ in this diatomic. The trend observed by increasing the number of ligands is identical for the three charge models and is in fact not surprising: q (Cl) always decrease with increasing n .

3.3 Spin-orbit coupling calculations

As already mentioned, the free Po⁴⁺ ion has a [Xe]4f¹⁴5d¹⁰6s² electronic structure. We recall here that within the non-relativistic or scalar-relativistic picture (*i.e.* when one considers orbitals and not spinors), the SOC is only active with the occurrence of partially filled p, d, f, *etc.* shells (note that this is a necessary but not sufficient condition). Therefore, electron donation to the Po⁴⁺ ion is required to activate the SOC, and in fact we have just seen that such a strong charge transfer occurs for the [PoCl]³⁺ complex, which encouraged us to analyse the role of the SOC on this Po–Cl bond, in particular concerning the bond distance.

At the non-relativistic or scalar-relativistic Hartree-Fock level, three bonding valence orbitals are doubly occupied and

have no populated antibonding counterparts (see Fig. 6). These are the doubly degenerate π -bonding HOMO and HOMO−1 orbitals and the HOMO−2 σ -bonding orbital. This σ -bonding orbital is lower in energy than the π -bonding ones, meaning that the molecular orbital diagram is uncorrelated. Below these, the two antibonding and bonding σ orbitals are essentially formed by linear combinations of the 6s orbital of Po and of the 3s orbital of Cl. Those two extra valence orbitals are doubly occupied, giving a net zero contribution to the formal bond order. Overall, the formal bond order in this system is 3, and the populated bonding orbitals are (i) delocalised on the two atoms (see Fig. 6) and (ii) essentially formed by linear combinations of the 6p orbitals of Po and of the 3p orbitals of Cl.

Without explicitly computing effective bond orders (EBOs),^{65–69} we would like to stress that two effects will tend to weaken the bond and thus enlarge the bond distance:

- Electron correlation may promote electron pairs from a bonding to an antibonding orbital, here especially from the σ orbital to the σ^* one (correlation induced reduction of the σ contribution to the EBO);
- The SOC may promote single electrons to antibonding orbitals, here especially the π orbitals to the σ^* one (SOC induced reduction of both the π and σ contributions to the EBO).

As a consequence, the effective bond order will significantly differ from the formal one, meaning that one should not see the Po–Cl bond in [PoCl]³⁺ as triple (and we have already seen with the delocalisation index, in the absence of the SOC, it is more likely to be a double bond).

In previous works,^{19,25} the role of the SOC on the molecular geometries of polonium(IV) complexes was neglected, based on reference to the electronic structure of the free Po⁴⁺ ion. However, since we have here evidenced that a strong effect

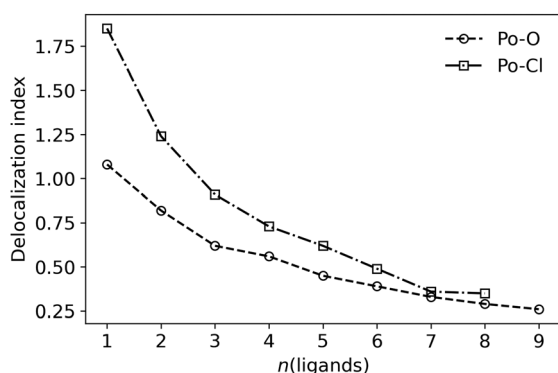


Fig. 5 Mean MP2/def2-TZVP Po–O and Po–Cl delocalisation indices as functions of the number of ligands (n) for polonium(IV) complexes with water molecules and chlorides, respectively.

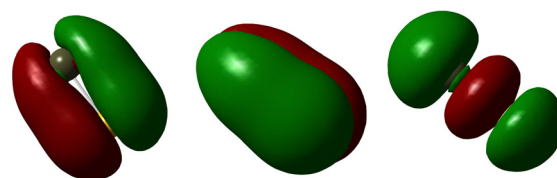


Fig. 6 Representations of the HOMO, HOMO−1 and HOMO−2 π and σ bonding orbitals of the [PoCl]³⁺ complex. Note that HOMO and HOMO−1 are degenerate by symmetry (those are the π orbitals).



can potentially occur in the hypothetical $[\text{PoCl}]^{3+}$ complex, we have computed molecular geometries in the absence and in the presence of the SOC, with two different frameworks, which are in fact complementary (see the Computational details section):

- ZORA scalar relativistic and two-component Hartree-Fock calculations (no electron correlation, *a priori* inclusion of the SOC).
- Two-step SOCI calculations, with a first scalar relativistic step and then diagonalisation of the $\mathbf{H}_{\text{tot}} = \mathbf{E}_{\text{el}} + \mathbf{H}_{\text{SOC}}$ matrix on the basis of the M_s components of the states of the previous step (spin-orbit free electron correlation, *a posteriori* inclusion of the SOC).

We just recall here that the first approach may overestimate the impact of the SOC (by lacking correlation) and that the second one (contracted SOCI) may underestimate it (by neglecting the SOC polarisation⁷⁰).

First, we tried out the *a priori* inclusion of the SOC, using the AMS software. For the $[\text{PoCl}]^{3+}$ complex, we observed an increase in the bond length from 2.17 to 2.29, *i.e.* a 0.12 Å SOC-induced bond lengthening. This is a significant effect, clearly larger than the expected error of the underlying quantum mechanical method (meaning that this effect is significant and thus not negligible). As we have seen that “covalency”, a key ingredient to activate the SOC in polonium(iv) complexes, quickly diminishes in the $[\text{PoCl}_n]^{4-n}$ series, we have performed calculations for the $n = 2, 3, 4$ and 8 complexes. For the $[\text{PoCl}_2]^{2-}$ complex, the SOC triggers a 0.01 Å bond lengthening, which becomes even more insignificant for larger n values. From this first SOC study, we can already conclude that the SOC is negligible in high-coordinated polonium(iv) complexes, implying that its inclusion should not be key to perform future calculations of polonium(iv) complexes in solution (solvent molecules will always surround the heavy cation). On the contrary, a significant SOC effect can potentially occur on low-coordinated polonium(iv) complexes, meaning that SOC should somehow generally be probed in diatomics or systems with only a couple of coordinated ligands (those systems are mostly of theoretical interest since gas phase experiments with polonium are not that safety-friendly).

Alternatively, we have also performed SOCI calculations on the $[\text{PoCl}]^{3+}$ complex, which proved to be interesting in terms of the SOC. We recall here that the two-step SOCI implementations do not yet propose any numerical or analytical gradients, meaning that the user has to explore the potential energy surfaces, here the potential energy curve (obviously with respect to the bond distance), by manual scanning.⁷¹ Several degrees of freedom are inherent to the SOCI approach: the size of the active space in the state-average complete active space (SA-CASSCF) calculation, the number of states that are included in the SA-CASSCF calculation and the way post-CASSCF electron correlation is introduced. In our calculations, we have considered 6 active electrons within 6 active orbitals (the σ , π , π^* and σ^* orbitals that are essentially formed by linear combinations of the 6p orbitals of Po and of the 3p orbitals of Cl). Note that test calculations showed that enlargement of the active space hardly affected our conclusions. With a ground state

spin-singlet state, only spin-triplet spin components are directly coupled to it. After some tests, we have set up the SA-CASSCF state space to 11 spin-triplet and 1 spin-singlet states. Post-CASSCF electron correlation has been introduced by replacing the E_{el} diagonal elements of the SOCI matrix^{72,73} by N-electron valence space perturbation theory at second order (NEVPT2) ones.^{74–76} With this setup, we observed a SOC-induced bond lengthening of 0.08 Å (from 2.27 to 2.35).

Before concluding on the role of the SOC on the bond distance, it is important to comment on the difference between both the scalar relativistic bond distances reported above (2.17 Å vs. 2.27 Å). For the sake of pedagogy and completeness, we would like to mention that three main factors may affect this difference: the basis set (TZ2P vs. a combination of DKH-def2-TZVP and SARC-DKH-TZVP, depending on the atom), the scalar relativistic Hamiltonian (ZORA vs. Douglas-Kroll-Hess) and the reference electronic structure method (Hartree-Fock vs. NEVPT2). As mentioned before, electron correlation is meant to weaken the bond order and in fact, also enlarge the bond distance. To probe this, we have performed an additional Hartree-Fock scan with ORCA with the same setup as in the second set of calculations, reproducing an equilibrium bond distance of 2.17 Å, as in the first set of calculations. We thus essentially attribute the observed difference in scalar relativistic bond distances to electron correlation.

Since the second approach is expected to underestimate the effect of SOC on the bond distance while the first one should overestimate it (see above), we finally conclude for a ~ 0.1 Å SOC-induced bond lengthening in $[\text{PoCl}]^{3+}$. This lengthening is of the same order of magnitude as the one observed for the AtX ($X = \text{F} - \text{At}$) diatomic molecule for instance,⁶⁹ for which the importance of the SOC was more obvious (the free At atom displaying a strong SOC splitting of ~ 2.9 eV⁶⁷ in its ground ^2P state). Also, it was shown in these systems that the SOC reduces the covalent character of the bond in terms of EBO⁶⁹ and of topological indicators^{59,77,78} such as the delocalisation index or the population of the bonding basin of the electron localization function.^{79–81}

4 Conclusions

In this work, two series of polonium(iv) complexes were investigated, respectively, with water and chloride ligands. A new benchmark study performed on the water complexes showed that the MP2/def2-TZVP level of theory is both accurate and efficient.

We have seen that the coordination sphere of the polonium(iv) ion may fit up to 9 water molecules or 8 chloride ligands, respectively. The interaction energy curves built by increasing n do not follow the exact same trend, a fact which we have attributed to the neutral vs. charged nature of the ligands.

We have also analysed the chemical bonds in the studied systems, generally qualified as regular closed-shell interactions. Because of signs of a strong “covalency” in the $[\text{PoCl}]^{3+}$



complex, we have investigated the influence of the SOC on the Po–Cl bond distance, quite unexpectedly revealing a SOC-induced bond lengthening of ~ 0.1 Å in this system, which vanishes for larger coordinations (in fact already starting from $n = 2$). This allows us to formulate a more general conclusion concerning heavy-element chemistry:

- Even if the reference free ion of a given heavy-element cation does not display any SOC, electron donation from the ligands to the cation may be strong enough to activate it and trigger sizeable effects. As a consequence, computational chemists should definitely not neglect the role of the SOC *a priori* (i.e. without any testing), especially in the case of low-coordinated complexes.

Two main perspectives arise from this work, (i) exploring more deeply and largely the role of the SOC in low-coordinated polonium(IV) complexes as well as with valence iso-electronic complexes such as bismuth(III) ones and (ii) studying thoroughly the effect of solvation on the molecular geometries, interaction energies and bonding in those complexes, which was out of the scope of the present work.

Author contributions

All the calculations were performed by N. Z. All the authors contributed to the scientific development of this work and jointly discussed the interpretation of the results. N. Z. wrote the first draft of the manuscript that has then been amended by all the co-authors.

Conflicts of interest

There are no conflicts of interest to declare.

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