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Properties of the Tetravalent Actinide Series in Aqueous Phase from a Microscopic Simulation Self-Consistent Engine[†]

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In the context of nuclear fuel recycling and environmental issues, the understanding of the properties of radio-elements with various approaches remains a challenge regarding their dangerousness. Moreover, experimentally some issues are also of importance; first, it is imperative to work at sufficiently high concentrations to reach the sensitivities of the analytical tools, however this condition often leads to precipitation for some of them, second stabilizing specific oxidation states of some actinides remains a challenge, thus making it difficult to extract general trends across the actinide series. Complementary to experiments, modeling can be used to unbiasedly probe the actinide's properties in aquatic environment and offers a predictive tool. We report the first molecular dynamics simulations based on homogeneously built force fields for the whole series of the tetravalent actinides in aqueous phase from Th^{IV} to Bk^{IV} and including Pu^{IV}. The force fields used to model the interactions among the constituents include polarization and charge donation microscopic effects. They are built from a self-consistent iterative quantum *ab initio* based engine that can be included in future developments as an element of a potential machine learning procedure devoted to generate accurate force fields. The comparison of our simulated hydrated actinide properties to available experimental data show the model robustness and the relevance of our parameter assignment engine. Moreover our simulated structural, dynamical and evolution of the hydration free energy data show that, apart from Am^{IV} and Cm^{IV}, the actinide properties change progressively along the series.

Because of its importance to separation science and to modeling actinide migration from nuclear waste repositories, there is a need of a profound knowledge of actinide environmental behavior. This involves a predictive understanding of actinide solution chemistry as the first critical step.¹ Namely, actinide environmental behavior implies speciation in the bulk phase but also at mineral-aqueous solution interfaces. For the later, sorption reactions can either involve pure electrostatic sticking on a

mineral surface with little change occurring in the first metal ion hydration shell, or inner-sphere surface complexation with chemical binding between the actinide and the mineral surface functional groups. Beside concentration, pH, and redox conditions, the competition between these two mechanisms is controlled by the stability of the cation first hydration shell, namely the hydration energy, as it governs its propensity to form strong inner sphere binding.²

In this context, the behavior of actinides, An^{IV}, in solution is highly relevant for two application fields connected to the nuclear industry : their migration as mentioned previously but also their separation from the spent nuclear fuels.³ However, the radiotoxicity and the complex chemistry of these elements explain the rather scarce availability of experimental data for them, in particular in aqueous phase. Whilst extended X-ray absorption fine structure (EXAFS) technique yields highly accurate An^{IV}/water oxygen bond lengths in the series Th⁴⁺-Bk⁴⁺, there are still large uncertainties in the experimental An^{IV} hydration numbers, N_c , ranging from eight to thirteen for Th⁴⁺-Bk⁴⁺.^{4,4-18}

To complement experimental data, research groups have either developed empirical theoretical models or performed quan-

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[†] Electronic Supplementary Information (ESI) available: They include

- Discussion of the accuracy of DFT for interaction energies
- Parameters of the force-field model for ion/water interactions
- Convergence of the BP-P process.

tum chemical computations on small An^{IV}/ligand clusters. We may refer here to the Born-like model (adjusted from geometrical experimental data) of David^{19,20} that yields An^{IV} hydration free energies, ΔG_{hyd} values. These values decrease regularly along the An^{IV} series. In 2003, David and Vokhmin²¹ revised the approach. They got then a very close tendency, at the remarkable exception of Np^{IV} whose ΔG_{hyd} value is down-shifted by more than 25 kcal mol⁻¹ with respect to the 1986 trend. A similar smoothly decreasing trend, although less steep, was also obtained by Bratsch and Lagowski.²²⁻²⁴ in 1986 with their simple model based on ionic radii. We may also cite the study of Banik et al.⁹ who performed quantum calculations on a set of small An^{IV} hydrated clusters (with bulk effects accounted for using a polarizable continuum solvent approach). These authors got An–O_{water} bond lengths in line with EXAFS data, and they concluded to a decrease of the An⁴⁺ hydration number from nine to eight, with a turning point at Cm⁴⁺.

To go beyond static models, molecular dynamic, MD, simulations have also been considered. However their predictive capability relies on the accuracy of the potential energy used to model microscopic interactions. Potential energies computed from quantum chemistry methods are the most relevant to investigate the properties of any kind of microscopic systems. Regarding An^{IV}, we may cite MD studies focusing on the hydration process of Th⁴⁺, U⁴⁺, and Pu⁴⁺ using density functional theory (DFT) methods.²⁵⁻²⁹ However none of the available DFT methods is able to describe accurately both water/water and cation/water interactions, at least when comparing DFT results to those of higher level quantum *ab initio* methods like the Møller-Plesset second-order perturbation theory, MP2, and above³⁰ (see discussions of Section 1 of the Supplementary Material). As high end quantum methods cannot be still considered to efficiently simulate bulk systems, even when used in hybrid quantum mechanical/molecular modeling schemes,^{31,32} molecular modeling approaches based on empirical interaction potentials, force fields, are still the most suited to simulate such systems on significantly long simulation times, and complex solution mixtures of different ions and different solvents.

As discussed earlier by other authors,^{30,33-35} An^{IV}/water microscopic interactions result from a complex interplay between mainly large electrostatic, non-additive polarization and charge-donation effects whose relative magnitude is not obvious to quantify. Moreover and as far as we know, most of the water force fields proposed to date are not able to capture the magnitude of the water/water repulsive interactions in cation first hydration shells, at the exception of the TCPE/2013 one.³⁵ This explains the difficulty of building well balanced force fields to study the hydration of An^{IV} and thus the very few An^{IV} MD bulk simulations based on force fields reported to date.^{25,30,35-37}

Besides the choice of a physically meaningful functional form for the force field, another critical issue is the strategy to assign force-field parameters. The most promising and robust strategy consists in assigning the parameters to reproduce only QM data regarding a limited set of training molecular clusters. These training data sets usually comprise small molecular clusters optimized in gas phase. We already built accordingly sophisticated force

fields, hereafter name Gas-Phase Parameter (GP-P), to study the hydration of Th^{IV} and Cm^{III}.^{30,37} However such an approach implicitly assumes that all the features of bulk phase interactions can be captured by investigating gas phase systems. In the particular case of U^{IV}, Atta-Fynn et al.²⁹ showed using quantum computations cation/water oxygen distances to be shifted by about 0.05 from gas phase to aqueous solution. Because of the +4 charge of An^{IV} elements, such a distance shift yields a change in the Coulombic interaction energy between An^{IV} and each water oxygen of its first hydration shell amounting to about 20 k_BT at ambient conditions. Hence a natural improvement, but more costly in term of computational resources, would be to consider bulk phase cluster structures for building accurate force field. We will denote this kind of force-field Bulk-Phase Parameter (BP-P). The generation of an increasingly larger set of reference data ensure the robustness of the force-field, and opens-up the possibility to handle complex mixtures of ions and solvents.

The most common parameter fitting process is a single step procedure consisting in best reproducing the properties of a usually restraint training molecular cluster set. The ongoing increase of the available computational resources yields research groups to propose more and more sophisticated fitting procedures where new QM target data are generated along the procedure cycles (see Figure 1 as an illustration), the latter can be used in machine-learning FF fitting process. We may refer here to Monte-Carlo sampling approach of Galbis et al.,³⁸ or the Force Balance method proposed by Wang et al.³⁹ Inspired by the available literature, we propose a coherent *ab initio* model associated to a self-consistent iterative procedure to homogeneously and coherently built a class of BP-P force fields for the whole An^{IV} series, in order to obtain metrical information on the coordination environment of the tetravalent actinide series from Th up to Bk, as well as to explore the evolution of the hydration free energies along the series, that will be confronted to the available empirical data. For comparison purpose, we also investigated the behavior of the lanthanide element Ce^{IV} that is often considered as a non radioactive surrogate for Pu^{IV}.^{40,41}

1 Methods and computational details

1.1 The force field

We use the rigid water model TCPE/2013.³⁵ For the ion/water interactions, we consider the *ab initio* based force field that describes the total interaction as the sum of four energy components,

$$\Delta U = U^{\text{rep}} + U^{\text{qq}'} + U^{\text{ct}} + U^{\text{pol}}. \quad (1)$$

The first three correspond to repulsive U^{rep} , Coulombic $U^{\text{qq}'}$ and charge-transfer U^{ct} contributions, the latter accounting for the partial "covalent" character of the actinide/water interactions. For a system of N atoms, the additive terms U^{rep} , $U^{\text{qq}'}$, and U^{ct} are defined as:

$$U^{\text{rep}} + U^{\text{qq}'} + U^{\text{ct}} = \sum_{i=1}^N \sum_{j,j>i}^N \left[A_{ij} \exp(B_{ij} r_{ij}) + \frac{q_i q_j}{4\pi\epsilon_0 r_{ij}} - D_{ij} \exp\left(-\frac{r_{ij}}{\beta_{ij}}\right) \right], \quad (2)$$

where r_{ij} is the distance between atoms i and j , $\{q_i\}$ are the static charges located on the atomic centers, and $(A_{ij}, B_{ij}, D_{ij}, \beta_{ij})$ are adjustable parameters.

Polarization effects are modeled using induced dipole moments \mathbf{p}_i expressed as

$$\mathbf{p}_i = \alpha_i \cdot \left(\mathbf{E}_i^q + \sum_{j=1}^{N_\mu} \mathbf{T}_{ij} \cdot \mathbf{p}_j \right). \quad (3)$$

All the N_μ nonhydrogen atoms are polarizable centers, i.e., a single point polarizability is located on each nonhydrogen atomic center. Their isotropic polarizability is α_i . \mathbf{E}_i^q is the electric field generated on the polarizable center i by the surrounding static charges q_j , and \mathbf{T}_{ij} is the dipolar interaction tensor. They both include short-range Thole's-like damping functions,^{30,35,42} with an adjustable damping parameter κ . All the ion-water force-field parameters optimized with the GP-P and BP-P procedures described in the coming section are listed in Table S1 in the ESI.

1.2 QM reference data for GP-P and BP-P force fields

The QM reference data sets will be composed of total binding energies and fragment interaction energies. The first ones correspond to the chemical reaction $\text{An}^{\text{IV}} + n\text{H}_2\text{O} \rightarrow \text{An}^{\text{IV}}(\text{H}_2\text{O})_n$ and are thus calculated according to the equation $BE = E_{\text{An}^{\text{IV}}(\text{H}_2\text{O})_n} - E_{\text{An}^{\text{IV}}} - nE_{\text{H}_2\text{O}}$. This energy is then corrected from the basis set superposition error between the actinide and the water molecules following the well established counterpoise correction method with the equation:

$$BSSE = E_{\text{An}^{\text{IV}}(\text{H}_2\text{O})_n}^{\text{An}^{\text{IV}}} - E_{\text{An}^{\text{IV}}}^{\text{An}^{\text{IV}}} + E_{(\text{H}_2\text{O})_n}^{\text{An}^{\text{IV}}(\text{H}_2\text{O})_n} - E_{(\text{H}_2\text{O})_n}^{\text{(H}_2\text{O})_n}, \quad (4)$$

where the subscript refers to the fragment considered and the superscript to the basis used. The corrected total binding energy is then:

$$BE^{cpc} = E_{\text{An}^{\text{IV}}(\text{H}_2\text{O})_n} - E_{\text{An}^{\text{IV}}}^{\text{An}^{\text{IV}}} - E_{(\text{H}_2\text{O})_n}^{\text{An}^{\text{IV}}(\text{H}_2\text{O})_n} + E_{(\text{H}_2\text{O})_n} - nE_{\text{H}_2\text{O}}, \quad (5)$$

where the superscript has been omitted when the basis set used is the fragment's own basis set. The fragment interaction energies are the counterpoise corrected interaction energies computed as:

$$E_{int}^{cpc} = E_{AB}^{AB} - E_A^{AB} - E_B^{AB}, \quad (6)$$

with the fragments A and B corresponding, on the one hand to a single water molecule of the $\text{An}^{\text{IV}}(\text{H}_2\text{O})_n$ cluster and on the other hand to the remaining atoms.

For all the terms defined above, using the Turbomole package (7.1 release)⁴³ we have performed RI-MP2 method^{44,45} for the closed-shell Th^{IV} or UMP2 for the open-shell systems (An^{IV} , $\text{An} = \text{Pa}$ to Bk), fixing the multiplicity of the computed state at the highest value. For each gas-phase aggregates, the energy calculations were preceded by an initial geometry optimization. For the actinides, the Stuttgart-Cologne "small core" relativistic effective core potential (60 e^-) were used in conjunction with the associated segmented basis set,⁴⁶⁻⁴⁸ while the augmented correlation consistent triple ζ basis sets of Dunning,^{49,50} namely the aug-cc-pVTZ, were used for the water molecules. The 1s core

electrons of the oxygen as well as the 5s, 5p, 5d electrons of the actinides were not correlated in the MP2 step.

1.3 Self-consistent BP-P force field procedure for An^{IV}

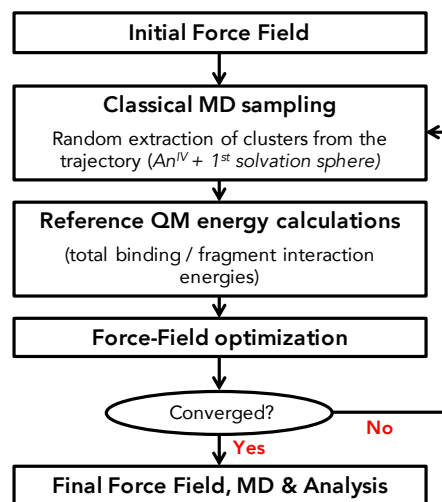


Fig. 1 Chart for the self-consistent iterative parameter optimization procedure. Each MD simulation is at the 10 ns scale. The sampling consists in extracting each 1 ps the An^{IV} plus its first hydration shell clusters. We select then 20 of these clusters whose MM binding energies, BE, corresponding to the reaction $\text{An}^{\text{IV}} + n\text{H}_2\text{O} \rightarrow \text{An}^{\text{IV}}(\text{H}_2\text{O})_n$, cover the range of the MM BEs corresponding to the full extracted cluster set. The reference QM computations are performed at the MP2 level.

The force-field parameters are assigned from the self-consistent iterative procedure, summarized in Figure 1. All iterative procedure iterations consist in the sequence of: MD sampling, QM reference energy calculation and fitting process. The parameter initial guesses correspond to force-field parameters assigned from the GP-P procedure, i.e. the training data set corresponds to small hydrated An^{IV} clusters comprising at most 10 water molecules optimized in gas phase using QM methods. The training data set at each iterative procedure's iteration comprises all the clusters from the previous two iterations, ensuring the BP-P convergence stability to the final force field. As high end QM computations regarding An^{IV} are still particularly computationally expensive, the MD sampling consists in extracting 20 hydrated An^{IV} clusters (the cation plus its first hydration sphere) from a 5 ns MD trajectory (See Section ??); for each cluster, about 10 interaction energies are computed to obtain the BEs and different fragment interaction energies. i.e 200 QM energies per cycle. To ensure that the 20 selected clusters at each iteration are representative, we checked that the MM BEs of these small clusters set span over the range of MM cluster BEs extracted from the full trajectory. At each iterative procedure's iteration, the evolution of the force-field parameters as well as structural properties (radial distribution function, and coordination number) is monitored, until the self-consistent solution is reached. Convergence is usually reached in less than 6 iterations, as illustrated in Figure S2 in the ESI. Today all the

computations of a single iteration of the procedure for the whole An^{IV} series can be performed at the week-scale using a few percents of a modern supercomputing system (made of about 100 000 computing cores).

The improvement in the force-field quality arising from the iterative procedure may be assessed from the data plotted in Figure 2. In this figure, the classical BEs computed from the parameter initial guesses (GP-P) and from the BP-P converged force field are plotted with respect to their QM reference counterparts for a training cluster set. If MM BEs from the initial parameter guesses are linearly correlated with QM ones, a large systematic deviation exists between the two BE sets, of about $-22 \text{ kcal mol}^{-1}$. If that represents only 2.5 % of the cluster BEs, that corresponds to about $36 k_B T$ at ambient conditions. With the BP-P force field, the BE mean error is centered on 0 and its standard deviation is reduced to less than 5 kcal mol^{-1} . These results support the earlier conclusion made by Tazi et al.,⁵¹ *i.e.* force-field parameter adjustments relying on realistic bulk-phase structures ensures the accuracy of the resulting force-field.

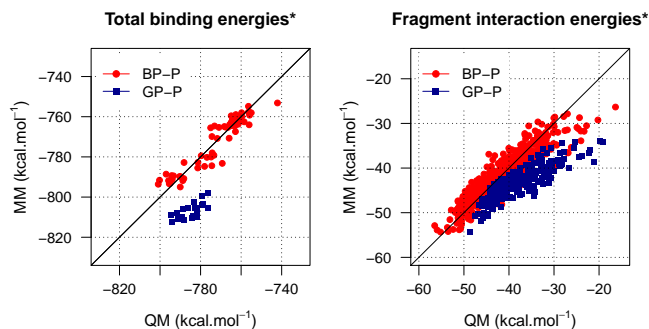


Fig. 2 Comparison of QM and MM binding energies BEs and of the fragment interaction energies, *i.e.* the interaction energy of a single water molecule of the $An^{IV}(H_2O)_n$ cluster with the remaining $An^{IV}(H_2O)_{n-1}$ fragment, for Th^{IV} when using the force-field parameter initial guess built from QM data corresponding to clusters optimized in gas phase (GP-P) and from the self-consistent iterative procedure (BP-P). The mean error is reduced to less than $0.1 \text{ kcal mol}^{-1}$ with a standard deviation of respectively 4.8 and $2.6 \text{ kcal mol}^{-1}$ for the total BEs and the fragment interaction energies with the latter approach.

1.4 Molecular dynamics simulations

The MD simulations of the An^{IV} in bulk water were conducted with a cubic box containing only one actinide cation for 1000 water molecules with periodic boundary conditions in PolarisMD package developed by one of us. The system is first equilibrated in volume and temperature in the $\{N,P,T\}$ ensemble with a Noose-Hoover thermostat and barostat.⁵² The different runs of production used for the parametrization are then run in the canonical ensemble⁵³ during 5 ns always preceded a 1 ns re-equilibration. The final step of the production from which thermodynamics data and structural informations are extracted, ran for 10 ns. It should be recalled here that the water model used for the simulations is the latest TCPE/2013 version.³⁵ The water molecule's structural parameters constrained to their bulk equilibrium val-

ues thanks to the RATTLE algorithm (the convergence criterium is set to $1 \times 10^{-6} \text{ \AA}$). Since the constituents of the system have no free internal motion, the simulation time-step is fixed to 1 fs and, in order to minimize the extra computational expense inherent to the polarization model, the induced dipoles are accounted for within the multiple time step r-RESPAP framework.⁵⁴ The hydration free energies, ΔG_{hyd} are computed relative to Th^{IV} using the standard Thermodynamic Integration (TI) scheme of twenty steps based on a linear interpolation of Hamiltonians. For each step, we performed a 100 ps run of equilibration followed by a 500 ps one during which the TI statistical average is computed.

2 Results and Discussion

The behavior of the whole An^{IV} series in 0.03 M cation aqueous solutions is studied by means of MD simulations at 10 ns scale using the BP-P force fields. We do not account explicitly for counter ions in our bulk simulations. However as the simulations are performed using an Ewald summation scheme with tinfoil boundary conditions, the cations are accompanied by uniform canceling background charge, acting as an implicit dilute counter ion cloud. We also simulated accordingly the lanthanide Ce^{IV} that is inferred to present very close hydration properties to Pu^{IV} . From MD trajectories, relevant structural properties like the pair An^{IV}/water oxygen radial distribution functions, RDFs, the position d_{AnO} of the RDF first peak, and the cation hydration number N_c are estimated.

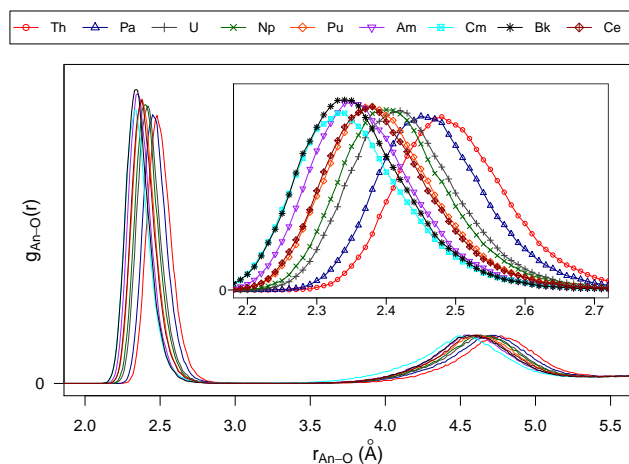


Fig. 3 An^{IV} – O pair radial distribution functions.

As expected from the An^{IV} large positive charge, MD simulations show the tetravalent cation surrounding water molecules to be organized in two well defined solvation shells (see Figure 3). At the exception of Cm^{IV} , the RDF first peak shifts in a very progressive fashion to shorter distances as the actinide gets heavier. The d_{AnO} distance decreases from Th^{IV} to Cm^{IV} , while the Bk^{IV} one is found to be slightly longer than for Cm^{IV} . Our d_{AnO} distance agree within at most 0.02 \AA compared to available EXAFS data (see Table 1). We plot on Figure 4 the d_{AnO} distances as a function of the An^{IV} ionic radii estimated from crystallographic data.^{20,55} The two series of data are linearly correlated. Follow-

ing the idea of Warren and Patel,⁵⁶ the ionic radii can be estimated from the distances d_{AnO} to which the water molecular radius is subtracted. For the TCPE/2013 model, the water radius is 1.393 Å.³⁵ That yields a nice agreement between our computed radii and the crystallographic ones, within less than 0.02 Å (see Table S3 in the ESI).

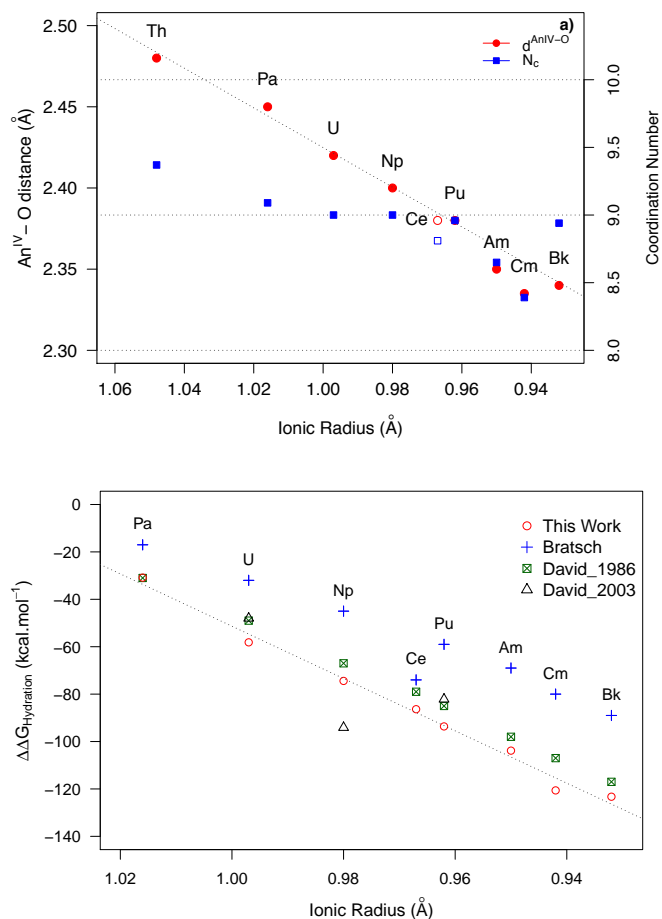


Fig. 4 (a) Distances d_{AnO} (red circles) as a function of ionic radii.^{20,55} The N_c estimates, computed by integrating the first RDF peak shown as blue squares in Figure 3, are reported on that plot (as the RDFs are null between their first and second peak the N_c error bar is zero). (b) An^{IV} hydration free energy relative to Th^{IV} ($\Delta\Delta G_{hyd}$) versus the An^{IV} ionic radii. In dashed lines, the linear regression fit, the regression coefficients are 0.99 and 0.99 for (a) and (b) respectively. Experimental data taken from the work by David et al.^{20,21} and Bratsch and Lagowski.²⁴

The computed N_c 's start at 9.4 for Th and decrease until they reach a plateau of about 9 for the series Pa–Pu, Bk with Am and Cm moving down away from the series with a coordination number of respectively 8.7 and 8.3, see Figure 4. Compared to experimental data regarding solutions for which the counter ions are known to not enter the An^{IV} first hydration shell (i.e. mainly perchlorate anions), N_c 's for the {Th^{IV} - Np^{IV}} series are included within the experimental range, while the N_c values for Pu^{IV} and Bk^{IV} are one unit above the experimental values. However, the EXAFS uncertainty regarding N_c values being 10% to 20%, our N_c values for Pu^{IV} and Bk^{IV} may be considered as agreeing with experiment, within the error bars.

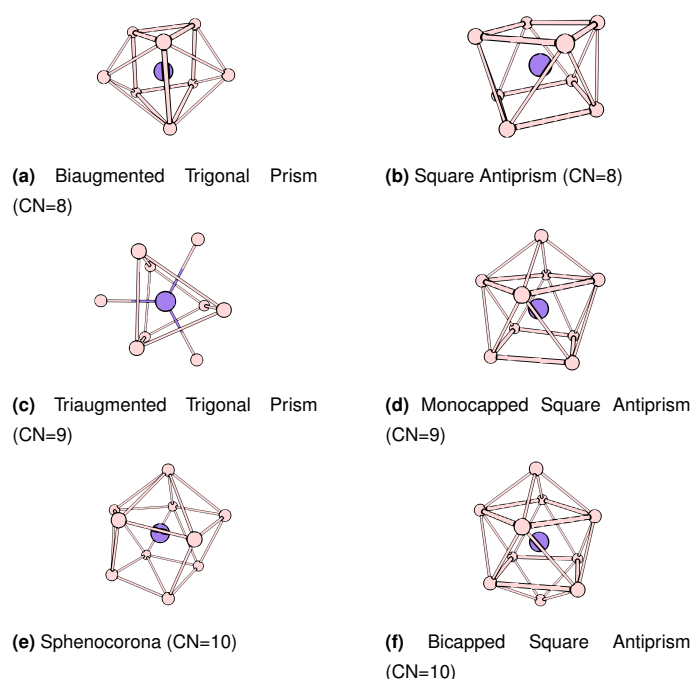


Fig. 5 Observed polyhedra for the coordination numbers (CN) ranging from 8 to 10.

The fractional N_c 's for Th^{IV}, Pa^{IV}, Am^{IV}, Cm^{IV} and Bk^{IV}, reveal the existence of an equilibrium between two coordination modes. An analysis of the coordination polyhedra has been made with *ChemNetworks*⁵⁹ and shows that the 8 coordination mode is associated with the formation of either a square anti-prismatic or a biaugmented trigonal prismatic geometry of the water molecules around the actinide cation. The identified polyhedra for the 9 coordination number are either a triaugmented trigonal prism or a mono-capped square anti-prism and for the 10 coordination number either a bicapped square anti-prism or a sphenocorona. The corresponding structures for the observed polyhedra extracted from the MD trajectories can be found in Figure 5. These coordination modes are in line with the ones observed for some studied An^{IV}, in particular for Th^{IV} and U^{IV}.^{26,28,29}

We computed the mean residence time, MRT, of a water molecule within An^{IV} first hydration shell along the MD trajectories from the approach of Impey et al.⁶⁰ The computed MRTs (See Table S2 in the ESI) range from a few hundreds of picoseconds for Cm^{IV}, Am^{IV} and Th^{IV} up to 1.5 ns for U^{IV}. Experimentally, only the U^{IV} MRT has been reported,⁶¹ about 185 ns, as well as an upper bound limit for Th^{IV}: 20 ns.⁶¹ Our estimates are *a priori* from one to two orders of magnitude smaller than experiment. This discrepancy may come from some deficiencies of our model to reproduce accurately the energy barriers in the water exchange reaction. On the other hand, we simulated here a single actinide cation surrounded by pure water whereas, in the experiment, very salty solutions are used (around 4 M of perchlorate). The presence of concentrated salts and counter-ions may also alter the cation hydration structure and affect the water dynamics at its vicinity.²⁹ The quality of the present BP-P force field cannot be incriminated yet to explain the apparent disagreement between the computed

Table 1 Comparison of structural MD computed data for the series of An^{IV}–water with GP-P and BP-P to experimental EXAFS data and other MD simulations.

Elt	GP-P		BP-P		Other MD simulations			Experiment		
	CN	d_{An-O} (Å)	CN	d_{An-O} (Å)	CN	d_{An-O} (Å)	Refs.	CN	d_{An-O} (Å)	Refs.
Th	10.0	2.48	9.4	2.48	8 - 11.4	2.40 - 2.50	25,29,30,36,37	9.0 - 12.7	2.44 - 2.46	4-8
Pa	10.0	2.44	9.1	2.45	-	-	-	-	2.43	9
U	9.9	2.41	9.0	2.42	8.7 - 9	2.45	26,57	8.7 - 10.6	2.40 - 2.42	4,10,11
Np	9.9	2.40	9.0	2.40	-	-	-	8.7 - 10.4	2.37 - 2.40	12-14
Pu	10.0	2.39	9.0	2.38	8.1	2.41	27	7.8 - 8.4	2.38 - 2.39	15,16
Am	9.0	2.33	8.7	2.35	-	-	-	-	-	-
Cm	8.9	2.32	8.3	2.335	-	-	-	-	-	-
Bk	9.5	2.34	8.9	2.34	-	-	-	7.9	2.32	17
Ce	9.1	2.365	8.8	2.38	9	2.44	58	< 9	2.42	18

and the rare experimental MRT estimates and further investigations should be carried on.

We computed the change in hydration free energies for the An^{IV} series relative to Th^{IV} ($\Delta\Delta G_{hyd}$) using the thermodynamic integration method. As for the distances d_{AnO} , we obtain a linear correlation between our $\Delta\Delta G_{hyd}$ values and the crystallographic ionic radii (see Figure 4). Our purely *ab initio* approach (BP-P force fields) predicts the $\Delta\Delta G_{hyd}$ values to smoothly decrease along the An^{IV} series. This is in line with the expectation that the continuous decrease of the d_{AnO} distances along that series yields a reinforcement of An^{IV}/water interactions. These results compare very well to David et al.'s estimates^{20,21} which all exhibit a decreasing trend much steeper than the one reported by Bratsch and Lagowski.²⁴ The discrepancies between the empirical models question the available data reliability and prove the challenges addressed to *ab initio* modelling approaches.

We also assigned BP-P parameters from the iterative procedure for the lanthanide element Ce^{IV}. We simulated then that element in an aqueous solution using the same computational protocol and simulation conditions as above. From our simulations, Ce^{IV} exhibits a very similar behavior to Pu^{IV} in bulk water, with an identical d_{AnO} distance of 2.38 Å and close N_c values, of respectively 9.0 and 8.8, for Pu^{IV} and Ce^{IV}. The similarity is further consolidated while comparing the respective RDFs of both elements that can be almost superimposed. Lastly, concerning the ΔG_{hyd} values, the Ce^{IV} estimate is found to be in between Np^{IV} and Pu^{IV}, still closer to the latter with a difference of 8 kcal mol⁻¹. Such a level of similarities between hydrated Ce^{IV} and Pu^{IV} have been also reported from QM and experimental studies.^{62,63}

Finally, considering the previous MD simulations reported in the literature, on the one hand, we find out that DFT-based MD generally underestimate the coordination number,²⁶ while exhibiting the largest computed interaction distances (Table 1); while on the other hand, different classical MD for Th^{IV} give very large ranges of coordination numbers (from 8 to 11.4) and interaction distances (from 2.40 Å to 2.48 Å). This disparity further proves the need for highly accurate force-field parameters assignment protocols to reduce as far as possible all the drawbacks mentioned in Introduction about building force fields.

3 Conclusion

In conclusion, we report the first polarizable An^{IV}–H₂O classical potentials derived from a self-consistent iterative *ab initio* methodology for the series from thorium to berkelium. We emphasize that the specific inclusion of sampled condensed phase reference structures in a self-consistent manner enables to reach an adequate accuracy necessary to simulate the actinide aqueous condensed phase systems. The hydration study from classical molecular dynamics with our newly developed force field reveals that, except for Am^{IV} and Cm^{IV} which sit aside the series, the structural and thermodynamical properties change in a very smooth fashion along the studied series. The first comparisons to experimental data exhibit a very good overall agreement. The increase of the hydration energies along the tetravalent actinide series has some impact on their geomobility. Indeed, one might expect all An(IV) to form like Th(IV) outer-sphere interactions with muscovite-like minerals,² while strongly complexing surfaces with carbonate, phosphate, or carboxylic acid may form inner-sphere complexes, calling for further extensions of force-field interaction models to such anionic ligands. Thus, this study will be extended in a near future by considering counter anions to achieve a thorougher comparability to experimental conditions. Finally, this results and methodological developments made here for the most highly charged ions of the periodic table are strictly transferrable to any ion or cation in solution for which an exploration of several properties in water or other solvent is envisaged, being either related to biological or atmospheric issues.

Conflicts of interest

There are no conflicts to declare.

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