Effect of Solvation on the Vertical Ionization Energy of Thymine: From Microhydration to Bulk

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Supporting Information

ABSTRACT: The effect of hydration on the vertical ionization energy (VIE) of thymine was characterized using equation-of-motion ionization potential coupled-cluster (EOM-IP-CCSD) and effective fragment potential (EFP) methods. We considered several microsolvated clusters as well as thymine solvated in bulk water. The VIE in bulk water was computed by averaging over solvent-solute configurations obtained from equilibrium molecular dynamics trajectories at 300 K. The effect of microsolvation was analyzed and contrasted against the combined effect of the first solvation shell in bulk water. Microsolvation reduces the ionization energy (IE) by about 0.1 eV per water molecule, while the first solvation shell increases the IE by 0.1 eV. The subsequent solvation lowers the IE, and the bulk value of the solvent-induced shift of thymine’s VIE is approximately −0.9 eV. The combined effect of the first solvation shell was explained in terms of specific solute−solvent interactions, which were investigated using model structures. The convergence of IE to the bulk value requires the hydration sphere of approximately 13.5 Å radius. The performance of the EOM-IP-CCSD/EFP scheme was benchmarked against full EOM-IP-CCSD using microhydrated structures. The errors were found to be less than 0.01⋯0.02 eV. The relative importance of the polarization and higher multipole moments in EFP model was also investigated.

1. INTRODUCTION

Ionization of nucleic acid bases (NABs) is relevant to radiation and photoinduced damage of DNA. In realistic environments, several types of interactions affect ionization energies (IEs) of the nucleic acid bases, quantities that govern charge migration, radiationless relaxation, proton transfer, and other photoinduced chemical changes and, therefore, ultimately control the redox properties and photostability of DNA. These include H-bonding interactions between the Watson−Crick pairs, π-stacking interactions between neighboring bases, interactions with sugar−phosphate backbone, counterions, and, of course, the omnipresent biological solvent, water.

Numerous gas-phase studies have characterized some of the above factors using model systems. For example, recent studies of NABs, their dimers, and microhydrated NABs illuminated the effects on different noncovalent interactions on IEs and hole delocalization patterns in ionized species. It was found that both π-stacking and H-bonding can reduce IEs by 0.4⋯1.0 eV via two distinctly different mechanisms. In π-stacked isomers, the IE reduction is due to the hole delocalization over the extended π-systems. In H-bonded species, large IE reduction is achieved via electrostatic stabilization of the localized hole by its interactions with the dipole moment of the favorably oriented “neutral” moiety. A similar mechanism is at play in microhydrated species. Microsolvation has been found to decrease the IEs of nucleobases by about 0.1 eV per water molecule; however, these findings cannot be extrapolated to bulk because, in addition to specific solvent−solute interactions, there are long-range contributions due to electrostatic screening and solvent polarization.

Solvent effects are expected to lower IEs of nucleobases and stabilize charged phosphate groups (thus, increasing their detachment energies), reducing the gap between nucleobases and phosphate IEs, suggesting that nucleotide ionization becomes more favorable in water. Ab initio calculations of isolated thymidine using CASPT2 have suggested that the preferred oxidation site in the nucleotide corresponds to the π orbital of the thymine base. This conclusion has been reassured by recent experimental study by Bradforth and co-workers who concluded that NABs are indeed preferred sites for ionization in DNA based on photoelectron measurements of vertical IEs of the pyrimidine nucleosides in aqueous microjets. This study reported that vertical ionization energy (VIE) of deoxythymidine in water is around 8.3 eV.
(i.e., red-shifted by 0.7—0.9 eV relative to gas phase). Based on DFT and continuum solvent model calculations, the authors concluded that the ionization occurs predominantly from the NAB and not the sugar or phosphate moiety.22

Due to inherent complexity of the problem causing experimental and computational challenges, there are still open questions regarding the effect of bulk solvation on IEs of the building blocks of DNA. For example, there are still no experimental measurements of IEs of solvated nucleobases. Most of the computational studies focused on either model microhydrated systems16,23–25 or employed implicit solvent models23,22,26–28 such as polarized continuum solvent (PCM) lacking the ability to describe specific solvent—solute interactions (e.g., g, h-bonding). Most of the PCM calculations focused on the adiabatic IEs, although the so-called nonequilibrium PCM has been used to calculate IEs.22 A recent study29 by Cau et al. has attempted to bridge this gap by modeling the effect of solvent and backbone environment on the lowest VIE of DNA bases by using a quantum mechanics/molecule mechanics (QM/MM) approach that includes explicit solvent—solute interactions (albeit empirically described). They have reported29 unphysically large blue shifts (3.2—3.3 eV) due to the environment, in disagreement with the experiment.22,30

This work presents calculation of the VIE of thymine in bulk water using explicit first-principle treatment of the solvent and high-level ab initio description of the solute. We employ a hybrid quantum mechanics/effective fragment potential (QM/EFP) approach, which is similar to QM/MM but does not rely on empirical force fields. The ab initio or QM part consists of the thymine molecule described by equation-of-motion ionization potential coupled-cluster with single and double substitutions (EOM-IP-CCSD).31–36 The water—thymine interactions are described by effective fragment potential (EFP) which includes Coulomb interactions with multipoles as high as octopoles obtained by Stones distributed multipole analysis45,46 and polarization interactions (see ref 40 for details on the EFP implementation employed in this study).

The effect of the environment (water) on the QM subsystem (thymine) is included via a perturbative technique (electronic embedding) in which the perturbed Hamiltonian of the QM system is

$$H = H_0 + \hat{V}$$

and the perturbation \(\hat{V}\) is given by

$$\hat{V} = \sum_{pq} \sum_A \left( \sum_k A_{kpq} \langle p | V | k \rangle + \sum_l A_{lq} \left( \frac{| Z_l |}{R_l} | q \rangle \right) \right)$$

where \(pq\) are the atomic orbitals in the ab initio region, \(A\) is an effective fragment, \(k\) denotes the multipole or polarization points depending on \(V_k\) and \(l\) denotes nuclei in the fragment and \(Z_l\) is the nuclear charge of the \(l\)th nucleus.

The induced dipoles of the effective fragments are iterated until self-consistency with each other and with the electronic wave function. Dispersion and exchange—repulsion interactions between the ab initio region and the effective fragments are treated similarly to the fragment—fragment interactions, that is, as additive corrections to the total energy. Thus, the total ground-state (or, more precisely, reference-state) energy of the QM/EFP system is

$$E_{GR}^{QM/EFP} = \langle \Phi_\text{gr} | H_0 + H_\text{Coul} + H_\text{disp} | \Phi_\text{gr} \rangle + E_\text{disp} + E_\text{rep}$$

where \(\Phi_\text{gr}\) is the reference-state wave function, \(H_\text{Coul}\) and \(H_\text{disp}\) are Coulomb and polarization EFP contributions to the Hamiltonian (subscript \"gr\" means that the induced dipoles corresponding to the electronic density of the reference state, which is often referred to as \"ground state\" , are used). \(E_\text{disp}\) is the electrostatic EFP-EFP energy; \(E_\text{rep}\) and \(E_\text{disp}\) include the exchange-repulsion and dispersion energies of both the EFP-EFP and the ab initio-EFP regions. \(E_{\text{pol}}\) is the self-consistent ground-state polarization energy of the QM/EFP system given by the following expression:

$$E_{\text{pol}} = -\frac{1}{2} \sum_{k} \sum_{a} \mu_{a}^{k} F_{a}^{\text{pol},k} + \frac{1}{2} \sum_{k} \sum_{a} \left( \frac{F_{a}^{\text{pol},k}}{a} \right) + \frac{1}{2} \sum_{k} \sum_{a} \left( \frac{F_{a}^{\text{pol},k}}{a} \right)$$
where $F^{\text{mult}}$ and $F^{\text{exc}}$ are the fields due to the static fragment multipoles and the nuclei of the quantum region; $F^0$ is the field due to the electronic density of the reference (Hartree–Fock, HF) state. $\mu^F$ and $\mu^k$ are the induced dipole and the conjugated induced dipole at the distributed polarizability point $k$. Note that the polarization contributions appear both in the quantum Hamiltonian through $\hat{H}^{\text{pol}}$:

$$\hat{H}^{\text{pol}} = -\frac{1}{2} \sum_k \sum_a \frac{\mu^k a}{R^3}$$

and in the EFP energy as $E^{\text{pol,EFP}}$.

The derivation and the programmable expressions of all EFP terms were discussed in the previous EFP papers. The details of the implementation used in this study can be found in ref 40.

In the EOM-IP-CCSD/EFP calculations, the reference-state CCSD equations for the cluster amplitudes $T$ are solved with the HF Hamiltonian modified by the electrostatic and polarization contributions due to the effective fragments, eq 1. The induced dipoles of the fragments are frozen in the coupled-cluster calculation. Such treatment, in which the dipoles are not reoptimized for the coupled-cluster wave function, is justified when the HF density provides a good zero-order approximation to the electron density of the coupled-cluster wave function and, consequently, the electric fields due to these densities are similar. For the systems considered in this work, the errors due to the neglect of the electron correlation contribution to the EF polarization were estimated to be less than 0.001 eV.

In EOM-CCSD, the target states energies, $E_k$, are found by diagonalizing the similarity-transformed Hamiltonian $\hat{H} \equiv e^{-T} \hat{H} e^T$:

$$\hat{H} R_k = E_k R_k$$

where $R_k$ is the excitation operator (with respect to the reference state $\Phi_0$), describing $k$th target state. In EOM-IP, the reference and target states have different numbers of electrons and the operator $R$ is of a Koopmans-like type.

The transformed Hamiltonian $\hat{H}$ includes Coulomb and polarization contributions from the EFP part. As $\hat{H}$ is diagonalized (using Davidson iterative procedure) in an EOM calculation, the induced dipoles of the effective fragments are frozen at their reference-state values, that is, the EOM equations are solved in a constant EFP field. To account for solvent response to electron rearrangement in the EOM target states (i.e., due to excitation or ionization), a perturbative noniterative correction is computed for each EOM root as follows. The one-electron density of each target EOM state (excited or ionized) is calculated and used to repolarize the environment, that is, to recalculate the induced dipoles of the EFP part in the field of this EOM state. These dipoles are used to compute the polarization energy corresponding to this state.

Following ref 39, the total energy of the ionized state with the inclusion of the perturbative response of the EFP polarization is

$$E^{\text{IP,QM/EFP}}_{\text{IP}} = E_{\text{IP}} + \Delta E_{\text{pol}}$$

where $E_{\text{IP}}$ is the energy found from eq 6 and $\Delta E_{\text{pol}}$ has the following form:

$$\Delta E_{\text{pol}} = \frac{1}{2} \sum_k \sum_a \left[ -\left( \mu^F_{ex,a} - \mu^k_{gr,a} \right) (F^{\text{mult}} a + F^{\text{exc}} a) 
+ \left( \mu^F_{ex,a} F^{\text{pol}} k a - \mu^k_{gr,a} F^{\text{pol}} a \right) 
- \left( \mu^F_{ex,a} - \mu^k_{gr,a} \right) \left( F^{\text{pol}} a + F^{\text{exc}} a \right) \right]$$

where $F^a_{\text{IP}}$ and $F^k_{\text{IP}}$ are the fields due to the reference (HF) state and the excited-state electronic densities, respectively; $\mu^F_{ex,a}$ and $\mu^k_{gr,a}$ are the induced dipole and conjugated induced dipole at the distributed polarizability point $k$ consistent with the reference-state density, whereas $\mu^F_{ex,a}$ and $\mu^k_{gr,a}$ are the induced dipoles corresponding to the excited state density.

The first two terms in eq 8 describe the difference of the polarization energy of the QM/EFP system in the ionized and ground electronic states; the last term is the leading correction to the interaction of the ground-state-optimized induced dipoles with the wave function of the excited state.

As discussed in detail in ref 39, the EOM states have both direct and indirect polarization contributions. The indirect term comes from the orbital relaxation of the solute in the field due to induced dipoles of the solvent. The direct term given by eq 8 is the response of the polarizable environment to the change in solute’s electronic density upon excitation. Unlike EOM-CCSD/EFP for the excitation energies, the direct polarization contribution in EOM-IP-CCSD/EFP can be very large (up to 0.7 eV for the systems considered in this study) since the electronic densities of the neutral and the ionized species are very different.

An important advantage of the perturbative EOM/EFP scheme described above is that it does not compromise multi-state nature of EOM and that the electronic wave functions of the target states remain (bi)orthogonal to each other because they are obtained with the same (reference-state) field of the polarizable environment. This would not be the case if polarization of the environment were treated fully self-consistently with the density of each electronic state.

3. COMPUTATIONAL DETAILS

The equilibrium geometries of microhydrated thymine clusters optimized by RI-MP2/cc-pVTZ are from ref 16 (the structures are also given in Supporting Information). These structures were used in the EOM-IP-CCSD calculations with the cc-pVTZ and 6-31+G(d) basis sets. The RI approximation was used in the former calculations.

In the QM/EFP calculations, the QM region consisted of the thymine molecule and was described by EOM-IP-CCSD/6-31+G(d). The water molecules were treated as effective fragments.

The EOM-IP-CCSD/EFP calculations were carried out using Q-Chem. The EF potentials of water are from the Q-Chem fragment library. These potentials were computed by using GAMESS following the protocol described in ref 40.

In the model systems, the structure of the thymine molecule was frozen at its equilibrium geometry (optimized by RI-MP2/cc-pVTZ) and the water molecule geometries are taken from the Q-Chem fragment library.

The equilibrium snapshots for thymine in bulk water were generated using molecular dynamics (MD) simulations carried out as follows. The AMBER-parm99SB/TIP3P solvent model was employed for the classical MD simulation of thymine and 1544 water molecules. For parm99SB, Restrained Electrostatic Potential (RESP) charges were calculated using the antechamber suite of AMBER9 following a geometry optimization at the B3LYP/cc-pVTZ level. Periodic boundary conditions were applied treating long-range electrostatic interactions with the particle-mesh Ewald technique. Prior to running production step, water box was allowed to relax. At this stage, position restraints were applied on all atoms of thymine. The following relaxation of the entire system was...
Table 1. VIEs (eV) of Microsolvated Thymine Computed Using the EOM-IP-CCSD/EFP, EOM-IP-CCSD/TIP3P, and Full EOM-IP-CCSD with the 6-31+G(d) Basis Set

<table>
<thead>
<tr>
<th>system</th>
<th>Full IP-CCSD</th>
<th>IP-CCSD/EFP</th>
<th>ΔIE (EFP)</th>
<th>IP-CCSD/TIP3P</th>
<th>ΔIE (TIP3P)</th>
</tr>
</thead>
<tbody>
<tr>
<td>T(H2O) (T1)</td>
<td>8.84</td>
<td>8.85</td>
<td>0.01</td>
<td>8.99</td>
<td>0.15</td>
</tr>
<tr>
<td>T(H2O) (T2)</td>
<td>8.88</td>
<td>8.90</td>
<td>0.02</td>
<td>8.89</td>
<td>0.01</td>
</tr>
<tr>
<td>T(H2O) (T3)</td>
<td>8.91</td>
<td>8.91</td>
<td>0.00</td>
<td>8.91</td>
<td>0.00</td>
</tr>
<tr>
<td>T(H2O)2 (T11)</td>
<td>8.72</td>
<td>8.72</td>
<td>0.00</td>
<td>8.84</td>
<td>0.12</td>
</tr>
</tbody>
</table>

Table 2. Effect of the Basis Set on VIE (eV) of Microsolvated Thymine Calculated Using Full EOM-IP-CCSD

<table>
<thead>
<tr>
<th>system</th>
<th>VIE (CCSD/cc-pVTZ)</th>
<th>∆VIE (CCSD/cc-pVTZ)</th>
<th>VIE (CCSD/6-31+G(d))</th>
<th>∆VIE (CCSD/6-31+G(d))</th>
</tr>
</thead>
<tbody>
<tr>
<td>thymine</td>
<td>9.14</td>
<td>n.a.</td>
<td>8.97</td>
<td>n.a.</td>
</tr>
<tr>
<td>T(H2O) (T1)</td>
<td>9.01</td>
<td>−0.13</td>
<td>8.84</td>
<td>−0.13</td>
</tr>
<tr>
<td>T(H2O) (T2)</td>
<td>9.05</td>
<td>−0.09</td>
<td>8.88</td>
<td>−0.09</td>
</tr>
<tr>
<td>T(H2O) (T3)</td>
<td>9.08</td>
<td>−0.06</td>
<td>8.91</td>
<td>−0.06</td>
</tr>
<tr>
<td>T(H2O)2 (T11)</td>
<td>8.89</td>
<td>−0.25</td>
<td>8.72</td>
<td>−0.25</td>
</tr>
</tbody>
</table>

4. RESULTS AND DISCUSSIONS

4.1. Benchmark Calculations of Thymine Microhydrates. Table 1 compares the values of the lowest VIE of microsolvated thymine computed by EOM-IP-CCSD/EFP and full EOM-IP-CCSD in the 6-31+G(d) basis set. The errors in IE's due to EFP approximation are less than 0.02 eV. Moreover, the correct ordering of the IE's in the different structures is preserved in the EFP calculations. Another important point is that the errors in EOM-IP-CCSD/EFP VIE's remain almost constant when going from one to two water molecules.

Table 2 demonstrates that the effect of the basis set on the solvent-induced shift in VIE (∆VIE) is negligible. However, the effect of the basis set on the absolute values of VIE's is considerable, that is, the difference between the cc-pVTZ and 6-31+G(d) values is about 0.17 eV. Thus, VIE of thymine in water can be reliably computed by combining solvent shifts (∆VIE) calculated with a smaller basis set and the VIE value of a bare thymine computed with a larger basis set. Using a smaller basis set in solvent-shift calculations allows us to perform extensive sampling over solvent configurations, which is important for converged results.

Figure 1 shows the results for higher VIE's. The EOM-IP-CCSD and EOM-IP-CCSD/EFP values agree well below 11.5 eV. The errors increase above 11.5 eV, as we approach the lowest VIE of water (≈11−13 eV66). As expected, the largest errors are observed for the states that have a significant electron density on the water molecule (see Figure 2). The state corresponding to removing the electron from the water moiety is simply missing in EOM-IP-CCSD/EFP. When the electron hole is delocalized over thymine and water, our separation of the system into the QM and EFP region becomes qualitatively incorrect; the QM/EFP approximation is valid as long as the orbitals (perturbed by the solvent) are localized on thymine. Fortunately, the gap between the IE's of the solute and the solvent provide a guideline for choosing QM/EFP (or QM/MM) separation; the ionized states are localized on the solute as long as they are considerably below the lowest IE of the solvent.65
4.2. Solvent Shifts Due to Specific Integrations. To understand the effect of individual hydrogen bonds at the different sites of thymine as well as the length over which the hydrogen-bond network affects VIE, we considered several model structures (Figure 3) with water molecules placed at different parts of the thymine molecule. Note that these structures do not represent equilibrium geometries of microhydrates (shown in Figure 1).

4.2.1. Effect of a Single Hydrogen Bond. In water–thymine hydrogen-bonding, the water molecules can act as donors or as acceptors. Figure 3 shows the positions of the water molecules with respect to thymine in the model structures constructed to probe the effect of different types of h-bonds on IE. When the water molecule is h-bonded to the NH groups of thymine, it acts as the electron donor and, therefore, reduces the VIE. Likewise, water that is hydrogen-bonded to the carbonyl groups acts as the electron acceptor, thereby increasing the VIE.

Table 3 shows the VIE shifts due to water molecules forming h-bonds with the CO and NH groups of thymine. We observe that a single water molecule h-bonded to a carbonyl group increases the VIE by \( \approx 0.29 \) eV, while a molecule h-bonded to a NH group reduces the VIE by \( \approx 0.34 \) eV. The magnitude of these opposing effects due to the different types of h-bonds provides a rationale for the observed VIE changes due to microhydration by a single water molecule. In the lowest-energy monohydrates (e.g., T1 structure shown in Figure 1), water is h-bonded to both the CO and the NH group, which yields the red shift of about 0.1 eV. Other microhydrates exhibit a similar trend of about 0.1 eV reduction in VIE per water molecule.16–18

The comparison between the VIEs calculated using EOM-IP-CCSD and EOM-IP-CCSD/EFP (see Table 3) further reinforces the excellent performance of the EFP treatment of water.

In sum, owing to their different nature, the h-bonds to CO and NH groups have opposing effects on VIEs. When water is bound to CO acting as electron acceptor, the VIE increases, whereas electron-donating water bound to NH leads to the VIE decrease. These effects of solvent–solute specific interactions cannot be described by continuum solvent models.

4.2.2. Length Scale of Water–Thymine Interactions. The h-bond network acts as conduits for the electrostatic interactions, thereby increasing or decreasing electron density on thymine and, consequently, affecting its IE. To understand the VIE of thymine in bulk water and to obtain a converged result, we need to determine the length scale over which water molecules can affect the VIE of thymine.

![Figure 2. MOs representing the ionized states of the T1 monohydrate computed by EOM-IP-CCSD and EOM-IP-CCSD/EFP. The errors in IEs due to the EFP approximation are also shown.](image)

![Figure 3. Model structures of thymine with one h-bonded water. Subfigures a,b,c,d refer to the structures with thymine and one water h-bonded to the CO group, whereas e and f refer to the structures with water h-bonded to the NH group.](image)

![Table 3. Effect of Specific h-Bonds on ΔVIE (eV) of Thymine Using Model Geometries Shown in Figure](image)

<table>
<thead>
<tr>
<th>ΔVIE</th>
<th>CO</th>
<th>NH</th>
</tr>
</thead>
<tbody>
<tr>
<td>method</td>
<td>a</td>
<td>b</td>
</tr>
<tr>
<td>full IP-CCSD</td>
<td>+0.27</td>
<td>+0.30</td>
</tr>
<tr>
<td>IP-CCSD/EFP</td>
<td>+0.27</td>
<td>+0.30</td>
</tr>
<tr>
<td>error</td>
<td>+0.00</td>
<td>+0.00</td>
</tr>
</tbody>
</table>

*The errors due to EFP treatment of water are also shown. The CO columns refer to structures a–d, whereas the NH columns refer to structures e and f.*

In liquid water, there are instantaneous h-bonds acting as electron density donors or acceptors. These competing effects cancel out quite efficiently obscuring the real length scale of
water-thymine interactions. Therefore, to determine the maximum distance over which water can affect thymine’s IE, we considered artificial chains of water molecules enhancing the electron donation/accepting effects. Four such model structures are shown in Figure 4. Model systems a and b are constructed by adding water molecules to the structure with water forming a single h-bond with the CO group. The subsequent waters are h-bonded to the O atom of the preceding water molecule (i.e., acting as h-donors). Model systems c and d are constructed by adding hydrogen-accepting water molecules to the structure with water forming a single h-bond with the NH group.

Figure 5 shows the convergence of VIE with respect to the number of water molecules in these structures. As one can see, the apparent length scale of the interactions is different in these structures. Structure b exhibits the slowest convergence. In general, the structures that have water molecule h-bonded to the NH group of thymine have shorter range of interaction than those in which water is h-bonded to the CO group. Figure 5 shows that the VIEs are affected by up to 9th and 20th water molecules in the case of water molecules h-bonded to the NH group, to be compared to 10th and 25th water molecules h-bonded to the CO group.

These results show that water molecules that are as far as 30 Å away from thymine can affect its VIEs. However, in a disordered system like liquid water, numerous instantaneous opposing interactions cancel out resulting in a reduced apparent interaction length.

4.3. VIE of Thymine in Bulk Water. 4.3.1. Convergence to the Bulk Limit with Respect to the System Size. The convergence of IEs (or detachment energies, DEs) with respect to the model system size has been shown to be quite slow. For example, the computed DE of iodide increases by about 2 eV when the first solvation shell of iodide (consisting of six water molecules) is encapsulated in a cluster of additional 858 waters.66 Even slower convergence of VDE with respect to the model system size was reported in ref 29 in which the convergence of VDE of aqueous chloride required using the box size of 30 Å, including approximately 1000 water molecules.

Figure 4. Different model structures used to study the length scale of water—thymine interactions.

Figure 5. Shift in VIE (ΔVIE, eV) as a function of number of water molecules in the model structures shown in Figure 4.
To estimate the minimum box size needed for converged VIE of solvated thymine we computed the VIEs averaged over multiple solvent configurations obtained from MD snapshots including different number of solvation shells.

We define solvation shells as follows. The first shell was defined to include water molecules whose H atoms are within 2.7 Å from the O atoms of thymine and those with O atoms that are within 2.7 Å of the H atoms of thymine. The position of the first minima in the radial distribution function between NH−O and CO−H. The position of the first minima are at 2.45, 2.65, 2.6, and 2.5 Å, respectively (as shown in the Supporting Information). Thus, the first shell was designed to include all water molecules that have a direct h-bond to thymine. The n-th shells are designed to include water molecules with the center of mass within $n \times 2.7$ Å from thymine. With these definitions, the first six hydration shells contain approximately 10, 50, 150, 300, 500, and 800 water molecules.

When this criterion and averaging the IEs over 25 MD snapshots are used, Figure 6 shows that the IE converges to the bulk value within the fifth shell. Thus, a simulation box of at least 30 Å ($2.7 \times 5 = 13.5$ Å radius around thymine) is required for the converged result. Such sphere contains about 500 molecules. This cutoff radius is very similar to the one observed in ref 29, contrary to our expectation that the IEs of the neutral species may be less sensitive to the solvent than DIgs of the anions, which exhibit much stronger solvent-induced shifts (several eV are common). Interestingly, the convergence is not monotonous; the first shell increases the IE, whereas the bulk value is lower than the gas-phase IE. The increase of the IE due to the interactions with the first shell is somewhat surprising, in view of the results for microhydrated species in which the IEs were shown to decrease with each water molecule. This finding is explained below.

**4.3.2. Effect of the First Hydration Shell on the VIE of Thymine.** Figure 7 shows the VIE shift due to the first solvation shell. The average $\Delta$VIE is about +0.09 eV relative to the gas-phase value. Taking into account small red shift ($-0.06$ eV) due to structural fluctuations of thymine, the solvent-induced shift is 0.15 eV. The increase in VIE can be rationalized by analyzing the instantaneous configurations of water molecules around thymine in the MD snapshots used for averaging. Figure 8 presents the analysis of instantaneous h-bonds formed at each snapshot. On average, there are 2.1 h-bonds with the CO group and 1.02 h-bonds with the NH moiety of thymine. The average increase in IE due to h-bond to CO $(+0.29$ eV) is smaller than the decrease in IE due to hydrogen-bond to NH $(-0.34$ eV); however, because there are more h-bonds to CO than to NH, we observe a net increase in VIE relative to the gas phase value. A simple estimate of the shift based on the average number of different h-bonds and assuming that the effects are additive is $+0.2$ eV, which is remarkably close to the observed value of $+0.15$ eV.

To summarize, the first hydration shell increases the VIE, in stark contrast to the observed VIE decrease in microhydrated clusters (section 4.1). This is because in the microhydrates, the water molecules form the maximum number of the strongest h-bonds with thymine to form the most stable structure. Typically, such structures are not representative of the true distribution of water molecules around the base, as in bulk water the interaction with other water molecules influences the positions of the water molecules in the first hydration shell. Moreover, the real structure of the first hydration shell at room temperature possesses not static (0 K global minima) but a statistic character. We observe that average coordination number for CO is larger than for NH (2.1 vs 0.9). Consequently, statistically there are more h-bonds with CO than with the NH group, which leads to the increase in VIE.

**4.3.3. Effect of the Outer Hydration Shells (2−6) on VIE.** The effect of the subsequent water shells is shown as a function of number of configurations in Figure 9. The cumulative averages of $\Delta$VIE were computed. Table 4 shows the average $\Delta$VIE and standard deviation for the increasing number of water shells. We notice that after the first hydration shell, all the subsequent shells lower the VIE. Although the convergence is achieved only around fourth or fifth shell, the effect of outer shells is relatively small compared to the first two shells and the VIE computed with the first and second shells is quite close to the bulk value.
The effect of the subsequent shells on VIE is more difficult to rationalize within a simple statistical picture, as was done for the first shell above, because the h-bond statistics become increasingly complicated. Electrostatic interactions, which provide leading contribution to the long-range water/solute interactions, are responsible for the observed VIE change due to third-sixth shells.

4.3.4. VIE of Thymine in Bulk Solution. The bulk VIE was calculated by averaging instantaneous VIEs over 100 configurations obtained from 10 ns trajectory at 300 K. All water molecules (∼1500) present in the simulation box (36 Å) are included in the EOM-IP-CCSD/EFP calculations.

Figure 9 shows ΔVIEs computed using 100 configurations from the MD snapshots. The average ΔVIE converges around 25–50 configurations. Thus, 100 configurations provide sufficient sampling. The converged solvent-induced VIE shift is −0.9 eV.

Using QM/MM, Cauè et al. reported that the combined effect of the environment (water and DNA backbone) on nucleobases leads to the VIE increase of about 3.4 eV. We observe an opposite effect due to solvation, possibly because we are considering only water environment. However, differences in methodology can contribute as well. Using the EOM-IP-CCSD/EFP/6-31+G(d) solvent shifts (ΔVIE) and the gas-phase VIE from the EOM-IP-CCSD/cc-pVTZ calculation, we obtain the VIE of bulk solvated thymine at ∼8.3 eV (9.2−0.9 eV). This is in excellent agreement with the experimentally observed VIE of nucleosides (8.3 eV). Figure 11 shows the distribution of the instantaneous ΔVIE of thymine at solvent configurations sampled in liquid water. The shifts are distributed from −1.75 to +0.25 eV. The most probable ΔVIE is −1.0 eV, which corresponds to the peak maximum in the photoelectron spectra of the first band. The lowest observed VIE corresponding to the onset of the first peak is ∼7.45 eV (9.2−1.75 eV).

Figure 12 shows the distributions of the instantaneous first and second VIEs of thymine in bulk water. We notice that there is some overlap between the two peaks (around 8.5–9.4 eV). The most probable VIEs corresponding to the first and second band maxima are 8.3 and 9.5 eV, respectively. The widths of the peaks due to solvation (full width at half maxima) are ∼0.75 and 1.0 eV, respectively, which is in excellent agreement with the experimental spectra.

4.4. Contributions of Various EFP Components to the VIE of Thymine. The change in VIEs of the thymine molecule due to water can be analyzed in terms of different components of the EFP potential, that is, Coulomb (charge, dipole, quadrupole, and octopole) and polarization energies. Such analysis is important for understanding the effects of hydration on the electronic structure of nucleobases.
for understanding the nature of water-thymine interactions. It is also of methodological significance as it may guide the development of new empirical water models and force fields.

The polarization component accounts for 23% of the fragment interaction energies owing to expected strong polarizability of extensive H-bonded systems. This is because polar molecules such as water and thymine can strongly perturb each other's wave functions and their interactions will result in significant charge redistribution. For example, our calculations show that the average dipole moment of thymine in water is 6.5 D, to be compared against the respective gas-phase value of 4.1 D. Likewise, the dipole moment of water in bulk water increases up to 2.9 D (the gas-phase value is 1.85 D).67 Our calculations also show that the dipole moment of water increases significantly even in microhydrates, for example, the values for thymine + 1H2O structures are 2.25, 2.45, and 2.35 D (in T1, T2, and T3), and in the dihydrate, the water dipole moment is 2.65 D. These changes in dipole moment quantify how strongly thymine polarizes the solvating water molecules.

Figure 13 shows the effect of polarization on the VIEs and the error due to the neglect of polarization. We observe errors as high as −0.8 eV (average error ≈ −0.6 eV). Thus, polarization is crucial for accurate calculations of VIEs in water.

The dipole, quadrupole, and octopole components account for 9, 12, and −0.2%, respectively, of the fragment interaction energies. The errors in the VIEs (see Figure 14) show similar trends due to the neglect of the dipole, quadrupole, and octopole moments, respectively. Thus, in this system, multipoles up to quadrupoles are of considerable importance. The effect of octopoles is small. Note that the multipoles used to carry out these benchmark calculations were not recomputed assuming the new truncation of multipoles (i.e., the multipoles are kept constant once calculated using up to octopole expansion).

Finally, for the benchmark purposes, we also employed the TIP3P point charges in calculations of VIEs of solvated and microsolvated thymine. Figure 15 shows the ΔVIE due to solvation in TIP3P water compared to the EFP waters and the differences between the respective VIEs. The average error (of the TIP3P VIE relative to EFP) is around 0.68 eV. About 80% of it is due to the neglect of polarization, and the remaining 20% is due to the neglect of higher multipoles.
multipole moments. Table 1 shows the VIEs of microsolvated water calculated with the TIP3P point charges. We notice that the errors can be as high as 0.15 eV in the case of the T1 structure.

To summarize, we conclude that point charges alone provide a poor description of the water molecules around thymine and higher multipole moments up to quadrupoles and polarization are necessary to accurately describe the VIEs.

5. CONCLUSION

We investigated the effect of microsolvation and bulk water on the lowest VIE of thymine using the EOM-IP-CCSD/EFP method. We benchmarked EOM-IP-CCSD/EFP against full EOM-IP-CCSD calculations for small systems \( [T(H_2O)]_n \ n = 1, 2 \). EOM-IP-CCSD/EFP values agree well with the full EOM-IP-CCSD calculations for VIEs below the lowest VIE of the solvent molecules (11–13 eV for various water clusters\(^{62–64}\)). For the lowest ionized states, the errors are below 0.02 eV. Using TIP3P point charges yields much larger errors even for small systems (about 0.15 eV for monohydrates), which increase with the system size approaching 0.7 eV for bulk water.

Calculations on the model systems demonstrated that the effect of water molecules on VIE depends on whether they form donor or acceptor \( H \)-bonds. Thus, a discrete QM/MM method is mandatory for a qualitatively correct description of such systems. The range of solvent–solute interactions and the effect of the various hydration shells also exhibits a delicate interplay of the specific \( H \)-bonds and long-range electrostatic/polarization effects. Our results allow us to estimate the smallest system required to understand the bulk effect of the water. We observed the convergence of VIE to the bulk value with respect to the system size by increasing the number of water molecules around thymine. Our data suggest that minimum 4–5 of water shells are needed, which corresponds to the approximately 30 Å simulation box.

We determined that the average value of VIE of thymine in bulk water is reduced by 0.9 eV. Thus, the predicted peak maximum of the lowest ionization band is 8.24 eV. Due to the solvent-induced broadening, the threshold value of the VIE is 7.39 eV. The computed second band maximum is at 9.4 eV. These values are in excellent agreement with the experimental VIE measurements.22 It was suggested that the phosphate moiety has little effect on the ionized states of nucleobases, possibly due to the screening effects. However, accurate calculations of solvated nucleotides (i.e., using EOM-IP-CCSD/EFP) are required to investigate this phenomenon.

A complete picture of the ionization process requires calculations of adiabatic IEs that involve solvation energy of the electron and reorganization energy of the solvent around the base. Furthermore, other bases should also be investigated, and the effect of the DNA backbone, interbase interactions, and counterions should be included. This is a subject of a future work.

ASSOCIATED CONTENT

Supporting Information. Additional data available. This material is available free of charge via the Internet at http://pubs.acs.org.

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(29) It should be noted that the experimental results are for the solvated nucleosides and not a solvated DNA molecule. It is conceivable that different interactions within DNA will result in different VIE.
(54) The effective fragments parameters were computed using Hartree–Fock wave functions. The exchange-repulsion, polarization and dispersion parameters were generated using the 6-31+G(3d,2p) basis set, whereas the electrostatic parameters were computed with 6-31+G(d) using Stone’s distributed multipole analysis. The mixed basis treatment of parameters is necessary for accurate analytical Coulomb multipoles, while keeping the bigger basis is required for the accurate description of the exchange-repulsion parameters.