Reactions of Alkaline Earth Metal Ions with Methanol Clusters

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The gas phase reactions between singly charged alkaline earth metal ions (Mg$^+$, Ca$^+$, Sr$^+$ and Ba$^+$) and methanol clusters are studied in a pick-up cluster source. In much the same way as for the alkaline earth metal ion–water cluster systems studied by Fuke et al. (J. Am. Chem. Soc. 1995, 117, 747), anomalous product distributions were observed that are characterized by two product switching regions. The first product switching from M$^+$(CH$_3$OH)$_n$ to MOCH$_3$$^+$$(CH$_3$OH)$_{n-1}$ occurs for n around 5. The MOCH$_3$$^+$$(CH$_3$OH)$_{n-1}$ species switches back to Mg$^+$(CH$_3$OH)$_n$ for n at around 15 (second switching). The critical sizes for the first and the second switching were found to be affected by the metal and deuterium substitutions. Ab initio calculations on the structures and energetics of the reactants and products were carried out to facilitate the interpretation of the product switching behaviors.

I. Introduction

The study of metal ion–solvent cluster (M$^+$–S$_n$) reactions provides valuable information on their reactivity patterns with increasing cluster size, reflecting the transition from gas-phase reactions to solution phase reactions. While research on the reactions of metal ions with molecules is abundant, studies on reactions of molecular clusters with metal ions have been scarce. In general, reactions of alkaline earth metal ions with solvent molecular clusters lead to the association products (e.g., Mg$^+$(CO$_2$)$_n$). However, unusual size-dependent reaction patterns may occur for some solvent clusters. Interesting examples are the reactions between Mg$^+$, Ca$^+$, and water clusters studied by Fuke’s group. The most intriguing finding from their experiments is that the association products M$^+$(H$_2$O)$_n$ (M = Mg and Ca) are unstable against H-elimination when the clusters reach a certain size. For example, Mg$^+$(H$_2$O)$_n$ change to MgOH$^+$(H$_2$O)$_{n-1}$ for n ≥ 6 (first switching). In addition, the H-elimination product switches back to Mg$^+$(H$_2$O)$_n$ for n ≥ 15 (second switching). Similar product distributions were observed for Ca$^+$–H$_2$O, though the exact switching points differ slightly.

To understand the reactions of M$^+$(H$_2$O)$_n$ (M = Mg and Ca), Iwata and co-workers performed ab initio molecular orbital calculations on these cluster ions and their corresponding dehydrogenation products, MOH$^+$(H$_2$O)$_{n-1}$, with n as large as 8 (for Ca$^+$). The two species were found to have very different structures. For instance, the first solvation shell of Mg$^+$(H$_2$O)$_n$ is filled with three water molecules and the fourth water molecule starts to fill the second shell. On the other hand, in MgOH$^+$(H$_2$O)$_n$, all the water molecules are directly bonded to Mg$^+$ for n ≥ 6. The hydration energy of MgOH$^+$ is much larger than that of Mg$^+$ because of the strongly polarized MgOH$^+$ core. As a consequence, the products in the reaction become more stable than the reactants with increasing cluster size. The energy change upon hydrogen elimination of Mg$^+$(H$_2$O)$_n$ is positive for n = 1–5, but becomes negative for n = 6, a result which is in good agreement with the observed first product switching.

These studies raised an interesting question as to whether such an anomalous cluster reaction would occur in other solvent clusters (e.g., CH$_3$OH). Investigation of other alkaline earth metal ion–solvent cluster systems may provide valuable information regarding the reaction mechanism. In particular, the substitution of one H in H$_2$O by a CH$_3$ group may help to understand the H-elimination behavior of M$^+$(H$_2$O)$_n$. On the basis of this consideration, we have systematically studied the reactions of four alkaline earth metal ions (Mg$^+$, Ca$^+$, Sr$^+$, and Ba$^+$) with methanol clusters using a pick-up technique. The substitution of H by a more bulky CH$_3$ group is expected to alter the structure of (H$_2$O)$_n$ substantially. In neat liquid, the nature of the hydrogen-bond structure is quite different for water and methanol. Water forms a three-dimensional, tetrahedral-based network such as ice, while methanol forms chains involving about six methanol molecules, on average. As such, the substitution of H by CH$_3$ offers an opportunity to investigate the structural effects of the molecular clusters on their reactions with alkaline earth metal ions. It would also be interesting to see whether H-elimination or CH$_3$-elimination is the dominant process.

There have been several reports regarding alkaline earth metal ion–methanol clusters with relatively small cluster size. Freisers’ group measured the bonding energy of Mg$^+$–methanol complex using Fourier transform mass spectrometry (FTMS) combined with the photodissociation technique, and obtained a value of 61 ± 5 kcal/mol. Theoretical calculations by Bauschlicher and co-workers on Mg$^+$(CH$_3$OH)$_n$ (n = 1 and 2) showed that Mg$^+$ ion is bonded to the oxygen atoms of the methanol molecules and that the bond is electrostatic in nature. The calculated binding energy for Mg$^+$–methanol is 41 ± 5 kcal/mol, which is smaller than the experimental result. It was argued that the value from the photodissociation experiment by Freisers et al. corresponds to an upper bound that is much larger than the true binding energy. Duncan and co-workers also performed photodissociation experiments on the Mg$^+$(CH$_3$OH)$_n$ complex. Unlike the earlier report by Freiser et al., who found Mg$^+$ to be the only ionic product, these authors observed several dissociation channels leading to
the formation of Mg\(^+\), CH\(_3\)\(^+\), MgO\(^+\), and MgOH\(^+\). The last product is difficult to ascertain because of the overlap of the isotope peaks of \(^{25}\)Mg and \(^{26}\)Mg in the mass spectrum. Farrar and co-workers studied Sr\(^+\)(CH\(_3\)OH) and Sr\(^+\)(CH\(_3\)OD) using the photodissociation method.\(^{12}\) They observed both the SrOH\(^+\) (SrOD\(^+\)) formation channel and the ligand loss channel leading to Sr\(^+\). Another dissociation channel corresponding to the SrOCH\(_3\)\(^+\) formation was not observed. For clusters as large as Sr\(^+\)(CH\(_3\)OH), the situation is essentially similar except that the large cluster size results in many possible dissociation products.\(^{13}\) It was found that dissociation to solvated SrOH\(^+\) yields precedence to the simple ligand loss channel with increasing cluster size.

Early studies concerning the reactions of alkaline earth metal ions with small methanol clusters have also been reported. Sato and co-workers studied the reaction of Mg\(^+\) with methanol clusters using the “laser ablation-molecular beam” method.\(^{14}\) Only the molecular addition products, however, were identified. In a flow-tube reaction studied by Castleman et al., successive solvation products were observed without dehydrogenation up to Mg\(^+\)(CH\(_3\)OH)\(_n\).\(^{15}\) Larger clusters were not produced in a significant amount. Very recently, Stace and co-workers reported the gas-phase chemistry of Mg\(^+\) and Mg\(^{2+}\) in association with methanol clusters using electron impact ionization.\(^{16}\) They observed ion signals corresponding to Mg\(^+\)(CH\(_3\)OH)\(_n\) and \([\text{Mg}(\text{CH}_3\text{OH})_n]^{2+}\) for \(n\) in the range 1–20. They also observed MgOCH\(_3\)\(^+\)(CH\(_3\)OH)\(_{n-1}\) starting at \(n = 3\) and ascribed it to the reaction products from Mg\(^+\)(CH\(_3\)OH)\(_n\). It should be pointed out that their experiment is quite different from ours as well as those of Fuke’s group in the sense that significant electronic excitation may be involved in the electron impact ionization process.

In this article, we present the results of our experimental and theoretical investigation of the alkaline metal ion—methanol cluster systems. We first describe the experimental setup and computational method. The experimental as well as computational results are then presented. Next, we discuss various aspects of the reactions between alkaline earth metal ions with methanol clusters with the focus on the first and second product switchings observed in the mass spectra. Finally, a summary is given in the end.

II. Experimental Methods

The experiments were performed using a pick-up cluster ion source coupled with a reflectron time-of-flight mass spectrometer (RTOFMS).\(^{17}\) Metal or metal oxides were mounted on a sample load lock, which was placed 9 mm downstream from a pulsed valve (R. M. Jordan) as shown in Figure 1a. The pulsed valve was used to generate the methanol cluster beam by supersonic expansion of a gas mixture of methanol vapor seeded in He with a back pressure of 7 atm through a 0.5 mm nozzle. The XeCl excimer laser beam was weakly focused on the sample with a flux of \(\sim 5 \times 10^9\) W/cm\(^2\). The laser-ablated metal ions perpendicularly crossed the expansion stream 10 mm from the ablation target where they reacted with the methanol clusters. The products were collimated with a 1 mm skimmer, and traveled 15 cm downstream to the extraction region of the RTOFMS. The product cations were extracted by a pulsed high voltage (950 V) and traveled along the field-free flight tube. The accelerated ions were reflected by a double-stage electric field at the end of the flight tube and detected by a microchannel plate detector (MCP). The source chamber and flight tube were differentially pumped by 10 and 6 in. diffusion pumps (Varian), respectively. The operating pressure was \(3 \times 10^{-6}\) Torr for the source chamber and \(2 \times 10^{-7}\) Torr for the flight tube.

The gas mixture of methanol—He was formed by bubbling the helium gas at a pressure of 7 atm through a methanol liquid reservoir. The pressure of methanol vapor in the gas mixture was estimated to be 127 Torr. An HPLC grade methanol sample (99.9% purity) was obtained from Fisher Scientific. For deuterium-substitution experiments, CH\(_3\)OD (Aldrich, +99.5% D), CD\(_3\)OH (RDH, 99% D), and CD\(_2\)OD (ACROS, 100% D) were used. All the samples were used without further purification. For the deuterated samples, a possible problem was the quick D/H exchange with the residual hydrogen species present in the nozzle system. Great care was taken when loading these samples to the reservoir to avoid contamination. The transfers of these samples were conducted under helium atmosphere in an AtmosBag (Aldrich).

A typical experiment started with the opening of the pulsed valve, followed by the firing of the ablation laser. The extraction high voltage was applied about 100 \(\mu\)s later. The delay times between the nozzle opening, the laser firing, and the ion extraction were adjusted to obtain the maximum signal intensities. The mass spectra were usually accumulated for 100 laser shots to increase the signal-to-noise ratio. We expect that the methanol cluster formation was largely quenched after expansion through the 8 mm distance (16 nozzle diameters) to the reaction region. Under a typical source condition, three-body collisions can be quenched in a few nozzle diameters for an axisymmetric expansion, while two-body collisions can remain in an extended length scale.\(^{18}\) Therefore, the reaction of metal cations with methanol clusters could be separated from the cluster formation process.

The cluster ion intensities were found to decrease quickly with increasing cluster size. To observe the large cluster ions, a pulsed high voltage was applied to two parallel metal plates.
inside the flight tube. These plates served as a mass gate to deflect unwanted ions. The mass gate was operated in such a way that, under normal conditions, the high voltage was always on except when the cluster ions of interest pass through the gate. A typical high-voltage setting for the mass gate was 800 V. By varying the pulse width and delay time, the mass spectra of different TOF regions could be obtained. Clusters containing as large as 25 methanol molecules were observed under typical experimental conditions.

We also measured kinetic energies of the laser-ablated metal ions. Figure 1b shows the setup for the ion kinetic energy measurements. The metal or metal oxide samples were attached to a sample mount on a load lock. The distance between the sample and the center of the ion extraction plates was adjusted to 15 cm. The laser energy and focusing conditions were kept the same as those in the reaction experiments. The pulsed laser irradiation of the sample generates a plume of neutral and ionic species. The ions then flew toward the ion extraction plates. When the ions traveled near the center of the ion extractor, a pulsed high voltage was applied to the plates. The ions were directed by the electric field along the flight tube and detected by the RTOFMS. To measure the kinetic energies of the metal ions, the delay time between the laser pulse and the ion extraction pulse was varied while monitoring the metal ion signal. The plot of the metal ion kinetic energy versus delay time gave the kinetic energy distribution of the laser-ablated metal ions.

III. Computation Methodology

Since a qualitative idea of the relative stability of the two species is of primary interest, we performed ab initio calculations at the SCF level with the electron correlation being ignored. This is probably reasonable, because an accurate description of the electrostatic interactions in the metal—ion system depends more on the completeness of the one-particle basis set than on the level of correlation treatment as pointed out by Bauschlicher et al.19 The geometries of the singly charged magnesium ion—methanol clusters, Mg+(CH₃OH)ₙ (n = 1—5), were optimized with the unrestricted Hartree–Fock self-consistent-field (UHF-SCF) method as Mg+ has an open-shell structure with a single 3s valence electron. The corresponding H-elimination cluster ions MgOCH₃+(CH₃OH)ₙ₋₁ (n = 1—5) were optimized with the closed-shell restricted Hartree–Fock self-consistent-field (RHF–SCF) methods. In all cases, the optimized geometries were obtained by varying all the geometric parameters.

The calculations were performed on a Silicon Graphics workstation using the Gaussian 94 package.20 The basis set 6-31G* was used. Natural charge distributions were obtained by performing natural bond orbital (NBO) calculation using the optimized geometry. Frequency calculations were also performed for the most stable isomers to obtain zero-point vibration (ZPV) energy corrections.

IV. Experimental Results

1. Metal Ion Kinetic Energy. Laser-ablation of magnesium metal and calcium oxides generates Mg⁺ and Ca⁺ as the predominant ion signals. Figure 2a shows the intensity of the Mg⁺ signal as a function of the delay time between the laser pulse and the ion extraction pulse at a laser fluence of 5 × 10⁵ W/cm². The Mg⁺ ions were detected at delay times between 9 μs and 27 μs. Since the travel distance is 15 cm, the velocities of these ions are estimated to be between 1.7 × 10⁶ and 5.6 × 10⁶ cm/s. The resulting kinetic energies of Mg⁺ range from 3.8 to 35 eV. The peak ion intensity is at a delay time of 18 μs based on the fitting. This corresponds to a velocity of 8.3 × 10⁵ cm/s and a kinetic energy of 8.6 eV.

Figure 2b shows the same plot for Ca⁺ at a laser fluence of 5 × 10⁵ W/cm². Most of the Ca⁺ signals were detected between the delay times of 18 μs and 41 μs. This translates to a velocity range from 8.3 × 10⁵ to 3.7 × 10⁶ cm/s and a kinetic energy range from 2.8 and 14.4 eV. The Gaussian fitting of the data reveals the existence of two kinds of Ca⁺ ions. The origin of the two peaks is unknown, but we note that this phenomenon was observed for other laser-ablated metal ions such as Cu⁺ from Y₁Ba₂Cu₄O₈ and was found to be related to the laser fluence.22 From the fitting in Figure 2b, the maximum Ca⁺ intensity is at a delay time of 32 μs, which corresponds to a velocity of 4.7 × 10⁵ cm/s and a kinetic energy of 4.6 eV. The measurements of kinetic energy distribution of Sr⁺ and Ba⁺ from laser ablation of SrO and BaO yielded similar results.

2. Reactions between Alkaline Earth Metal Ions and Methanol Clusters. 2.1 Reaction between Mg⁺ and Methanol Clusters. Typical TOF mass spectra of the reaction products between Mg⁺ and methanol clusters are displayed in Figure 3. In the mass spectrum in Figure 3a, Mg⁺ and its complexes with methanol clusters can be readily identified due to the characteristic isotope distribution of Mg (²⁵Mg, 78.99%; ²⁶Mg, 10.00%; and ²⁷Mg, 11.01%).23 Hereafter, the Mg we refer to is the most abundant ²⁵Mg isotope. Presumably, the cluster ions Mg⁺(CH₃OH)ₙ stem from the association reaction between Mg⁺ and methanol clusters:

\[
\text{Mg}^+ + (\text{CH}_3\text{OH})_m \rightarrow [\text{Mg}^+\text{(CH}_3\text{OH})_m]^+ \\
\rightarrow \text{Mg}^+\text{(CH}_3\text{OH})_n + (m-n)\text{CH}_3\text{OH}
\]  

The association products are probably stabilized by evaporation of one methanol molecule or by collision with the carrier gas. For clusters larger than Mg⁺(CH₃OH)_₅, the cluster ion
intensity decreases rapidly with increasing cluster size. To observe the large cluster ions, a pulsed high-voltage deflector was used to deflect the most abundant small cluster ions so that the detector gain can be increased. By appropriately choosing the pulse delay, pulse length, amplitude of the pulsed high voltage, and the voltage of the MCP detector, we could detect large cluster ions in a relatively broad TOF window.

The mass spectrum of larger cluster ions after reducing the small cluster ion intensities by the pulsed high voltage deflector is shown in Figure 3b. Clusters with up to 20 methanol units can be clearly observed. While it may appear at first glance that the mass peaks could be fitted to the formula $\text{Mg}^+(\text{CH}_3\text{OH})_n$, careful mass analysis of the spectrum reveals that the H-elimination products $\text{MgOCH}_3^+(\text{CH}_3\text{OH})_{n-1}$ were produced with a characteristic size distribution. The $\text{Mg}^+(\text{CH}_3\text{OH})_n$ cluster ions dominate the products for small cluster ions until $n = 5$. At $n = 5$, a significant intensity dip in the mass spectrum in Figure 3 marks a sharp switching of the reaction product from $\text{Mg}^+(\text{CH}_3\text{OH})_5$ to $\text{MgOCH}_3^+(\text{CH}_3\text{OH})_4$ through the following reaction:

$$\text{Mg}^+(\text{CH}_3\text{OH})_n \rightarrow \text{MgOCH}_3^+(\text{CH}_3\text{OH})_{n-1} + \text{H}, \quad n \geq 5$$  (2)

This product switching can be more clearly seen in Figure 4. For $n = 4$, only a single large peak corresponding to $\text{Mg}^+(\text{CH}_3\text{OH})_4$ dominates along with two small peaks due to the isotopes of the Mg. At $n = 5$, however, two relatively large peaks appeared, corresponding to $\text{MgOCH}_3^+(\text{CH}_3\text{OH})_4$ and $\text{Mg}^+(\text{CH}_3\text{OH})_5$, respectively. Here the intensity of the former is only slightly larger than that of the latter. However, for $n > 5$, $\text{MgOCH}_3^+(\text{CH}_3\text{OH})_{n-1}$ becomes the dominating species with very little $\text{Mg}^+(\text{CH}_3\text{OH})_n$ series left. This reaction pattern continues with increasing cluster size until $n = 15$, where $\text{Mg}^+(\text{CH}_3\text{OH})_n$ becomes the major species again as seen in Figure 5. This constitutes the second reaction pattern switching for the metal ion-solvent cluster system.

Further analysis of the mass spectra shown in Figures 3–5 revealed that there are some other reaction products with much smaller intensities. The first corresponds to protonated methanol clusters $\text{H}^+(\text{CH}_3\text{OH})_n$ (labeled as $A_n$). We suspect that these species are from an electron impact ionization process, during...
which the needed electrons were provided by the laser ablation process. The intensities of this series of cluster ions are rather small and quickly decrease with increasing cluster size. For clusters with \( n > 9 \), the cluster ion cannot be observed.

Another series labeled as B, starts to emerge for \( n \geq 5 \) as seen in Figure 4. Interestingly, it corresponds to the \( \text{MgOH}^+(\text{CH}_3\text{OH})_{n-1} \) series. This series of cluster ions was not observed for small \( n \) and starts at a similar cluster size as \( \text{MgOCH}_3^+(\text{CH}_3\text{OH})_{n-1} \). We attribute this series to the product of the following reaction with \( \text{CH}_3 \) elimination:

\[
\text{Mg}^+/(\text{CH}_3\text{OH})_n \rightarrow \text{MgOH}^+/(\text{CH}_3\text{OH})_{n-1} + \text{CH}_3
\]  

(3)

The intracluster reaction of \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) appears to have two elimination channels: H-elimination and CH3-elimination, with the former being the predominating process. The thermodynamic data shows that \( \text{CH}_3-\text{OH} \) bond dissociation energy is 92 kcal/mol, a value that is smaller than that of the \( \text{CH}_3\text{O} - \text{H} \) bond (105 kcal/mol).23 The facile elimination of \( \text{H} \) is perhaps due to its small size and the isotopic distribution of its \( s \) electron,24 which makes it kinetically more favorable than the \( \text{CH}_3 \)-elimination. The \( \text{MgOH}^+/(\text{CH}_3\text{OH})_{n-1} \) cluster ion coexists with \( \text{MgOCH}_3^+/(\text{CH}_3\text{OH})_{n-1} \) for \( n = 5 - 15 \). However, for \( n \geq 15 \), both cluster ion intensities decrease quickly and switch to cluster ions with one mass number difference. \( \text{MgOCH}_3^+/(\text{CH}_3\text{OH})_{n-1} \) shifts back to \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) while \( \text{MgOH}^+/(\text{CH}_3\text{OH})_{n-1} \) shifts to a cluster ion series with a formula \( \text{Mg}_2\text{O}_n\text{H}^+(\text{CH}_3\text{OH})_n \) (labeled as \( D_n \)).

It is also interesting to note another series \( \text{Mg}_2(\text{OCH}_3)_3^+/(\text{CH}_3\text{OH})_{n-2} \) (labeled as \( C_n \) in Figure 4), originating from the reaction of magnesium metal dimer cation \( \text{Mg}_2^+ \) with the methanol clusters. In this case, two hydrogen atoms have been eliminated and the eliminated product may very well be \( \text{H}_2 \) simply from the energetic arguments. A surprising finding in this series of cluster ions is that they can further dehydrogenate to give \( \text{Mg}_2(\text{OCH}_3)_2^+/(\text{CH}_3\text{OH})_{n-3} \) (labeled as \( E_n \)).

2.2. Deuterium Substitution Effects. To investigate the isotope effect for the reactions described above, experiments were performed for \( \text{Mg}^+ \) with deuterated methanol samples, namely \( \text{CH}_3\text{OD} \), \( \text{CD}_3\text{OH} \), and \( \text{CD}_2\text{OD} \). These isotope substitution experiments also serve to confirm our previous mass assignments.

The mass spectra for the reactions of \( \text{Mg}^+ \) with \( (\text{CH}_3\text{OH})_n \), \( (\text{CH}_3\text{OD})_n \), \( (\text{CD}_3\text{OH})_n \), and \( (\text{CD}_2\text{OD})_n \) are presented in Figures 6 and 7. To facilitate comparison, the x-axis in the mass spectra is converted to the number of methanol molecules \( n \) in the cluster. Evidently, the mass resolution of the spectra is sufficiently high to resolve one unit of difference in mass for clusters of as large as 16 \( \text{CD}_2\text{OD} \) molecules.

The reaction products for clusters containing 4–7 methanol molecules can be identified in the mass spectra in Figure 6. This region corresponds to the first product switching, where the \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) cluster ions dehydrogenate to give \( \text{MgOMe}^+/(\text{CH}_3\text{OH})_{n-1} \). For \( \text{CH}_3\text{OH} \), this switching occurs at \( n = 5 \) as described above. When the OH group in \( \text{CH}_3\text{OH} \) is substituted by the OD group, the switching point is clearly shifted to larger clusters by one methanol unit. For both \( \text{CH}_3\text{OD} \) and \( \text{CD}_2\text{OD} \), the association product ions \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) are still dominant for \( n \geq 5 \), although small peaks corresponding to \( \text{MgOMe}^+/(\text{CH}_3\text{OH})_{n-1} \) start to appear on the left sides of the strong peaks for \( n = 4 \) and 5. Only for \( n \geq 6 \) do the series \( \text{MgOMe}^+/(\text{CH}_3\text{OH})_{n-1} \) really take over the series of the association product ions. For \( \text{CD}_2\text{OH} \), the situation is very similar to that of \( \text{CH}_3\text{OH} \). The first product switching occurs at \( n = 5 \) where the intensity of \( \text{MgOCD}_3^+/(\text{CD}_3\text{OH})_4 \) becomes larger than that of \( \text{Mg}^+/(\text{CD}_3\text{OH})_5 \). This similarity is not surprising since the dehydrogenation reaction involves only the \( \text{O} - \text{H} \) bond cleavage, on which the replacement of \( \text{CH}_3 \) with \( \text{CD}_3 \) has little effect.

Shown in Figure 7 are the mass spectra of the product ions for clusters containing 13–16 methanol molecules. A second product switching from \( \text{MgOMe}^+/(\text{CH}_3\text{OH})_{n-1} \) back to \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) is clearly seen in Figure 7. The mass resolution in this high-mass region is slightly lower, but is still high enough to distinguish the difference of one atomic mass unit. Again, the x-axis of the mass spectra in Figure 7 is converted to the \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) cluster size \( n \) for the sake of comparison.

For \( \text{CH}_3\text{OH} \) and \( \text{CD}_2\text{OH} \), the critical sizes for the second switching are the same (i.e., both are at \( n = 15 \)). For example, in the intracluster reactions of \( \text{Mg}^+/(\text{CD}_2\text{OH})_n \), the intensities of the product ions \( \text{MgOCD}_3^+/(\text{CD}_3\text{OH})_{n-1} \) at \( n = 14 \) are very strong, and there are almost no \( \text{Mg}^+/(\text{CD}_2\text{OH})_{14} \) signals. However, at \( n = 15 \), the H-elimination product appears as a shoulder on the left of the stronger \( \text{Mg}^+/(\text{CD}_2\text{OH})_{15} \) peak. The intensities of \( \text{MgOCD}_3^+/(\text{CD}_3\text{OH})_{n-1} \) ions decrease further with increasing cluster size. In much the same way as seen in the first product switching, it appears that the \( \text{CH}_3/\text{CD}_3 \) exchange has no effect on the critical size of the second product switching.

The effect of OH/OD exchange on the second product switching was also examined. In contrast to the situation for the \( \text{CH}_3/\text{CD}_3 \) exchange, where there was no observable effect, the OH/OD exchange shifts the second product switching point to smaller clusters by one methanol unit. As can be appreciated in Figure 7, for both \( \text{CH}_3\text{OD} \) and \( \text{CD}_2\text{OD} \), the \( \text{MgOMe}^+/(\text{CH}_3\text{OH})_{n-1} \) ion intensity is larger than that of \( \text{Mg}^+/(\text{CH}_3\text{OH})_n \) at \( n = 13 \), but their intensities are reversed at \( n = 14 \).
The reactions of Ca$^{+}$, Sr$^{+}$, and Ba$^{+}$ with methanol monomer and dimer.

The H-elimination reaction leads to the breakage of the O–H bond in CH$_3$OH and this is compensated by the formation of M$^+$–OCH$_3$ bond. The binding energies of M$^+$–OCH$_3$ are not available yet. However, considering that CH$_3$ is more polarizable than H and that this would increase the ligand contribution to the bonding, we expect that the binding energy for M$^+$–
OCH₃ should be larger than that of M⁺−OH. Thus, although the CH₂O−H bond dissociation energy is 105 kcal/mol,²³ which is 13 kcal/mol larger than that of CH₃−OH, the large bond dissociation energy of M⁺−OCH₃ can still make the H-elimination reaction exothermic for Ca⁺, Sr⁺, and Ba⁺. Especially for Ba⁺, very few association products Ba⁺(CH₃OH) are formed compared to BaOH⁺ and BaOCH₃⁺, possibly due to the larger binding energies for Ba⁺−OH and Ba⁺−OCH₃. Again, for Mg⁺, the H-elimination reaction is still endothermic.

The results presented above also indicate that the reaction between the laser-ablated alkaline earth metal ion and methanol molecule occurs on the ground-state surface. The ⁵P excited state of Mg⁺ is known to be 102.2 kcal/mol above the ground state ⁵S.²⁸ Such a high internal energy is far more than enough to induce the CH₃-elimination and H-elimination reactions with CH₃OH, leading to the formation of MgOH⁺ and MgOCH₃⁺. However, the fact that these two products were not observed implies that the excited-state metal ion species did not participate in the reaction.

Another important point in Figure 8 is that the reactions of the alkaline earth metal ions with methanol dimer give only the association products. This is surprising especially for Ca⁺, Sr⁺, and Ba⁺ since, based on thermodynamic data, the CH₃-elimination and H-elimination reactions should be more exothermic for the methanol dimer than for the monomer. It is likely that, for the methanol dimer reaction, the collision complex can be stabilized by evaporation of the constituent methanol molecules, yielding the association product ion without reaction. Another possibility is that the lifetime of the collision complex is so long that it is eventually stabilized by the collision, most probably with the carrier gas, again forming the association product ion. This continues to be the case until the cluster size reaches ~5, where the first product switching takes place.

The mass spectra of the product ions from Ca⁺(CH₃OH)ₙ, Sr⁺(CH₃OH)ₙ, and Ba⁺(CH₃OH)ₙ and Mg⁺(CH₃OH)ₙ in the cluster size range from n = 3−7 are compared in Figure 9. In general, in this size range, the first product switching takes place for all the four alkaline earth metal ions although the exact cluster size for product switching may be somewhat different. As we have shown above, Mg⁺(CH₃OH)ₙ starts to change to MgOCH₃⁺(CH₃OH)ₙ₋₁ at n = 5. For Ca⁺, this critical size changes to n = 4. As can be seen from the spectra, for n = 3, the Ca⁺(CH₃OH)₃ ion still dominates, although a small peak corresponding to CaOCH₃⁺(CH₃OH)₂ already appears on the left of the strong peak. Right at n = 4, their intensities are reversed. This reaction product series continues for even larger clusters up to the second product switching region (see Figure 7) at n = 15, where CaOCH₃⁺(CH₃OH)ₙ₋₁ shifts back to Ca⁺(CH₃OH)ₙ. This critical size for second product switching is the same as that of Mg⁺.

Similarly, the first product switching for the reactions of Sr⁺ and Ba⁺ takes place at n = 5 and n = 6, respectively, as can be seen from the mass spectra in Figure 9. Although the mass resolution has deteriorated to a certain extent for larger clusters, careful analysis of the mass spectra reveals that the critical size of the second product switching for the reactions of Sr⁺ is 15. For the reactions of Ba⁺, at least for n ≤ 12, BaOCH₃⁺(CH₃OH)ₙ₋₁ are still the dominant cluster ions. The small signal intensity prevents further analysis for even larger clusters of Ba⁺(CH₃OH)ₙ with respect to the second product switching.

In general, the reactions of Ca⁺, Sr⁺ and Ba⁺ with methanol clusters (CH₃OH)ₙ (n > 1) exhibit a characteristic product distribution similar to that of Mg⁺ in terms of the first and second product switchings. The overall results are presented in Table 1.

**TABLE 1: Product Distributions for Reactions of Alkaline Earth Metal Ions with Methanol Clusters**

<table>
<thead>
<tr>
<th>reactants</th>
<th>metal ion</th>
<th>methanol</th>
<th>products</th>
</tr>
</thead>
<tbody>
<tr>
<td>(M⁺)</td>
<td>CH₃OH</td>
<td>M⁺(CH₃OH)ₙ</td>
<td>MΟMe⁺(CH₃OH)ₙ₋₁</td>
</tr>
<tr>
<td>Mg⁺</td>
<td>CH₃OH</td>
<td>n ≤ 4 and n ≥ 15</td>
<td>5 ≤ n ≤ 14</td>
</tr>
<tr>
<td></td>
<td>CH₃OD</td>
<td>n ≤ 5 and n ≥ 14</td>
<td>6 ≤ n ≤ 13</td>
</tr>
<tr>
<td></td>
<td>CD₃OH</td>
<td>n ≤ 4 and n ≥ 15</td>
<td>5 ≤ n ≤ 14</td>
</tr>
<tr>
<td></td>
<td>CD₃OD</td>
<td>n ≤ 5 and n ≥ 14</td>
<td>6 ≤ n ≤ 13</td>
</tr>
<tr>
<td>Ca⁺</td>
<td>CH₃OH</td>
<td>n ≤ 3 and n ≥ 15</td>
<td>n = 1 and 4 ≤ n ≤ 14</td>
</tr>
<tr>
<td>Sr⁺</td>
<td>CH₃OH</td>
<td>n ≤ 4 and n ≥ 15</td>
<td>n = 1 and 5 ≤ n ≤ 14</td>
</tr>
<tr>
<td>Ba⁺</td>
<td>CH₃OH</td>
<td>n ≤ 5 *</td>
<td>n = 1 and 6 ≤ n ≤ 12</td>
</tr>
</tbody>
</table>

* Signals are too weak for n ≥ 12.

V. Computational Results. We performed ab initio calculations on singly charged metal ion–solvent clusters Mg⁺(CH₃OH)ₙ and their corresponding H-elimination products MgOCH₃⁺(CH₃OH)ₙ₋₁. The size of the cluster ions in the present study is limited to n = 5 by the available computation capability. Methanol is a quite large subunit compared with water, and thus the ab initio calculations become very difficult for clusters containing a large number of methanol molecules. Even for a medium-sized cluster ion, many possible isomers may exist originating from the different orientations of the neighboring subunits. We do not intend to perform a comprehensive theoretical study to locate all the possible isomers at high levels. Instead, it is sufficient for our purpose to get a qualitative picture of the gross geometries and relative thermodynamic stability of the two species.
TABLE 2: Calculated Total Energies of Different Structures for Mg\(^{+}\)(CH\(_3\)OH\(_n\)) (n \(\leq 5\)) at the SCF/6-31G* Level as Well as the Relative Energies for Different Isomers

<table>
<thead>
<tr>
<th>species</th>
<th>structure</th>
<th>energies (hartree)</th>
<th>relative energies (kJ/mol)</th>
<th>ZPE (hartree)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mg(^{+})</td>
<td>1</td>
<td>-199.352 158 045</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>CH(_3)OH</td>
<td></td>
<td>-115.034 237 279</td>
<td>0.055 307</td>
<td></td>
</tr>
<tr>
<td>Mg(^{+})(CH(_3)OH)(_2))</td>
<td>2a</td>
<td>-429.532 100 589</td>
<td>0.115 055</td>
<td></td>
</tr>
<tr>
<td>Mg(^{+})(CH(_3)OH)(_3))</td>
<td>3a</td>
<td>-544.600 618 244</td>
<td>0.172 370</td>
<td></td>
</tr>
<tr>
<td>Mg(^{+})(CH(_3)OH)(_4))</td>
<td>4a</td>
<td>-659.668 910 660</td>
<td>0.231 335</td>
<td></td>
</tr>
<tr>
<td>Mg(^{+})(CH(_3)OH)(_5))</td>
<td>5</td>
<td>-774.728 233 616</td>
<td>0.290 120</td>
<td></td>
</tr>
</tbody>
</table>

*ZPE represents the zero-point energy corrections to the total energies for the most stable isomers.

For Mg\(^{+}\)(CH\(_3\)OH)\(_2\)), two isomers were studied, which result from the different orientations of the two OH groups in the two methanol molecules. For both structures, the two methanol molecules were found to be located on the same side of Mg\(^{+}\) as there is a high electron density on the other side of Mg\(^{+}\). The interaction with solvent molecules results in a large polarization of the 3s electron density away from the ligands. As shown in Figure 10, for structure 2a, the hydrogen atom in one of the OH groups points in the opposite direction from the hydrogen atom of the OH group in another methanol molecule. In structure 2b, the two OH groups in the two methanol molecules are close to each other, and the steric effect makes the two hydrogen atoms point away from the plane containing the Mg\(^{+}\) and the two oxygen atoms. This is similar to the situation in Mg\(^{+}\)(H\(_2\)O)\(_2\)), where the two water molecules are staggered. Despite the large steric hindrance expected for structure 2b, the two structures were found to be very close in energy. The structure 2b is found to be only 0.21 kJ/mol less stable than structure 2a. And the O–Mg–O angles in the two structures are also very close; both are around 92\(^\circ\), which is close to that in Mg\(^{+}\)(H\(_2\)O)\(_2\)).

Bauchlicher et al. used a larger basis set TZ2P to calculate the structures of Mg\(^{+}\)(CH\(_3\)OH)\(_n\)) for n = 1 and 2.\(^{10,11}\) Basically, the structures they found are very similar to what we obtained at the SCF/6-31G* level. There is only a slight difference in the Mg–O distances and the O–Mg–O angle. For example, we found the Mg–O distance in Mg\(^{+}\)(CH\(_3\)OH) to be 2.028 Å, and the calculation by Bauchlicher et al. gave a value of 2.01 Å.\(^{10}\) The difference in the O–Mg–O angle is found to be even smaller.

For Mg\(^{+}\)(CH\(_3\)OH)\(_3\)), two structures were investigated as shown in Figure 11. In structure 3a, all the three oxygen atoms in the methanol molecules are directly bonded to Mg\(^{+}\). Whereas, in structure 3b, two methanol molecules are directly bonded to Mg\(^{+}\), forming the first solvation shell, and the third methanol is bonded to the two methanol molecules in the first shell through two hydrogen bonds. Essentially, the three methanol molecules form a six-membered ring with all the six atoms in the ring within the same plane. Our calculation at the SCF/6-31G* level shows that structure 3a is more stable than structure 3b, although the energy difference between these two isomers is not significant. The relative energy difference is calculated to be 6.87 kJ/mol. This is in contrast to that of the Mg\(^{+}\)–water cluster system where the energy difference between the two corresponding isomers of Mg\(^{+}\)(H\(_2\)O)\(_3\)) cluster was calculated to be 14.4 kJ/mol at the SCF/6-31G level.\(^{5}\) In all probability, the larger size of methanol molecules has some effect on the relative stability between structures 3a, b. The larger ligand–ligand repulsions could make structure 3a less favorable as opposed to Mg\(^{+}\)(H\(_2\)O)\(_3\)) with a similar structure. However, the most stable isomer for Mg\(^{+}\)(CH\(_3\)OH)\(_3\)) is still the one with three methanol molecules directly bonded to Mg\(^{+}\) to form the first solvation shell.

Three isomers, whose structures are shown in Figure 12, were investigated for Mg\(^{+}\)(CH\(_3\)OH)\(_4\)). Similar to the situation of the
smaller clusters, due to the high electron density located on one side of Mg$^+$, all the solvent methanol molecules in the clusters prefer to be located on the other side of Mg$^+$. Structures 4a,b both have three methanol molecules in the first solvation shell, and the fourth methanol molecule starts to fill the second solvation shell. However, there are some differences between structures 4a,b. In structure 4a, the fourth methanol molecule is bonded to two methanol molecules in the first shell through two hydrogen bonds. They form a six-membered ring. For structure 4b, the fourth methanol molecule is bonded to only one methanol molecule in the first shell through one hydrogen bond. The six-membered ring structure in structure 4a is very similar to that in isomer 3b for Mg$^+$(CH$_3$OH)$_3$. With such a six-membered ring structure containing two hydrogen bonds, the isomer 4a is expected to be more stable than the isomer 4b with only one hydrogen bond. Our calculation at the SCF/6-31G* level shows that structure 4a is indeed more stable than structure 4b with an energy difference of 10.73 kJ/mol. But the hydrogen-bond length in structure 4b (1.768 Å) is much smaller than that in structure 4a (1.940 Å). Thus the number of hydrogen bonds is the major factor in determining the relative stability of the isomers. Presumably, the longer hydrogen bonds in structure 4a result from the large ligand–ligand repulsion of the two methanol molecules in the first shell, both of which are bonded to the fourth methanol molecule. Structure 4c corresponds to a structure with all the four methanol molecules directly bonded to Mg$^+$, forming the first solvation shell. This structure is found to be 17.01 kJ/mol higher in energy than structure 4a, obviously due to its larger ligand–ligand repulsions among the methanol molecules in the first solvation shell. Thus our calculation shows that the first solvation shell of Mg$^+$(CH$_3$OH)$_n$ is completed with three methanol molecules, and the fourth methanol molecule will start to fill the second solvation shell.

The methanol molecules in the second shell are connected to the methanol molecules in the first shell through hydrogen bonds with their neighboring OH groups. Again, this situation is quite similar to that of Mg$^+$(H$_2$O)$_n$ in which it was found that the most stable isomers also have three solvent molecules in the first solvation shell and the addition of more solvent molecules will start the filling of the second shell.

It is interesting to examine more closely the effect of second-shell formation on the solvent molecules in the first shell. The three methanol molecules in the first shell, which are all directly bonded to Mg$^+$ in structure 4a, can be divided into two types. The first type refers to the two methanol molecules, which are both connected with the methanol molecule in the second shell through hydrogen bonds. The second type then embodies the only methanol molecule, which does not bond directly with the methanol molecule in the second shell. From our calculation, it was found that the Mg–O bond distance for the first type of methanol molecules (2.091 Å) is slightly smaller than that for the second type (2.095 Å). This indicates that the interactions of the methanol molecules in the first shell with the methanol molecules in the second shell result in small geometry change of the methanol molecules of the first shell. The second obvious change is the natural charge of the oxygen atoms and hydrogen atoms in the OH groups of methanol molecules. The charge of oxygen atoms in the first type of methanol molecules is $-0.946$, which is more negative than that in the second type methanol molecule which is $-0.923$. The charge of hydrogen

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**Figure 11.** Optimized structures of Mg$^+$(CH$_3$OH)$_3$ (3a, 3b) at the SCF/6-31G* level. Angles are in degrees and distances are in angstroms. The values in square brackets indicate the natural charge for the corresponding atoms.

**Figure 12.** Optimized structures of Mg$^+$(CH$_3$OH)$_4$ (4a, 4b, 4c) at the SCF/6-31G* level. Distances are in angstroms. The values in square brackets indicate the natural charge for the corresponding atoms.
atoms in the OH groups of the first type of methanol molecules is found to be 0.543, which is more positive than that in the second type of methanol molecule by 0.02. In other words, the OH groups are more polarized by the hydrogen bonds with the methanol molecules in the second shell. This phenomenon was observed for all Mg\(^{+} (\text{CH}_3\text{OH})_5\) cluster ions having the second solvation shell. For example, in structure 4b, the charges of O and H atoms in the OH group, which is hydrogen bonded with the methanol in the second shell, are −0.968 and 0.562, respectively. This results from the strong hydrogen bond with a bond length of only 1.768 Å, which is much smaller than that in structure 4a. The bond length of Mg−O (2.073 Å) connected to the first type of methanol molecule is also significantly shorter than those of the other two Mg−O bonds (∼2.10 Å).

For Mg\(^{+} (\text{CH}_3\text{OH})_5\), the largest cluster we investigated, the fifth methanol molecule is expected to be in the second shell as well and bonded to the methanol molecules in the first shell via hydrogen bonds. Figure 13 shows the optimized structure of Mg\(^{+} (\text{CH}_3\text{OH})_5\). There are three methanol molecules in the first shell, which are all directly bonded to Mg\(^{+}\). The fourth methanol molecule is in the second shell and is bonded to two methanol molecules in the first shell through two hydrogen bonds similar in a way to that for the isomer 4a of Mg\(^{+} (\text{CH}_2\text{OH})_5\). They form a six-membered ring. The fifth methanol molecule is bonded to two methanol molecules; one is from the first shell, which is not hydrogen bonded to the fourth methanol molecule, and the other is from the second shell. This forms an eight-membered ring, which deforms the planarity of the six-membered ring. Such a stable structure can be considered as being derived from structure 4a. One would wonder if certain stable structures could also be derived from the isomers 4b,c. The isomers derived from 4b can have two structures. In the first structure, the fifth methanol molecule is bonded to the two idle methanol molecules in the first shell and further bonded to the fourth methanol molecule through hydrogen bonds. This will form two types of rings, a structure which is essentially the same as structure 5 in Figure 13. The second structure requires the fifth methanol molecule to be bonded to the fourth methanol molecule and one idle methanol molecule in the first shell. This can only form one ring and is not expected to be as stable as isomer 5. Similarly, the structure derived from 4c can also form only one ring, and the large ligand–ligand repulsions among the four methanol molecules in the first shell make it even more unstable. We therefore believe that the structure 5 corresponds to the most stable isomer for Mg\(^{+} (\text{CH}_3\text{OH})_5\).
The distances between Mg$^{2+}$ and the oxygen atoms in methanol are slightly larger than those in smaller clusters due to the increased ligand–ligand repulsion, but they are still smaller than those in Mg$^{n+}$ (CH$_3$OH)$_n$.

For larger clusters (e.g., MgOCH$_3^+$ (CH$_3$OH)$_4$), the question arises again as to whether the second solvation shell has started to form as the ligand–ligand repulsions increase. We investigated two isomers for MgOCH$_3^+$ (CH$_3$OH)$_4$ shown in Figure 15. Structure Va has all the five oxygen atoms directly bonded to Mg, and the methanol molecules are all located in the first solvation shell. The five oxygen atoms form a nearly trigonal-bipyramid structure commonly observed in five-coordination systems. The oxygen atom in MgOCH$_3^+$ is located at the vertex of the triangle that bridges the two pyramid structures. This oxygen atom forms a hydrogen bond with a hydroxide group in a methanol molecule nearby. The distance for this hydrogen bond is calculated to be 2.038 Å. The distances between this oxygen atom and other OH groups are all above 2.5 Å and do not form any hydrogen bond. The Mg–O distance increases further from that in MgOCH$_3^+$ (CH$_3$OH)$_3$. For example, the Mg–O distance in the MgOCH$_3^+$ core of MgOCH$_3^+$ (CH$_3$OH)$_3$ is 1.841 Å, which is much larger than that in MgOCH$_3^+$ (CH$_3$OH)$_4$ (1.774 Å). Obviously the ligand–ligand repulsions in structure Va are much larger than those in smaller cluster systems. Structure Vb has three methanol molecules directly bonded to MgOCH$_3^+$, so there are four oxygen atoms in the first solvation shell. The fifth methanol molecule is located in the second shell and it is connected to two methanol molecules in the first shell via two hydrogen bonds. The length of the hydrogen bond is 1.953 Å. With these two hydrogen bonds, a six-membered ring structure was formed to stabilize the system. Despite this stabilization, we found that this structure is higher in energy than structure Va by 11.52 kJ/mol. This is ascribed to the stronger electrostatic interactions between Mg$^{2+}$ and the oxygen atoms in structure Va than those in structure Vb.

We did not proceed the calculations further for even larger clusters, but it is expected that the stable structure for MgOCH$_3^+$ (CH$_3$OH)$_5$ may still have all the methanol molecules in the first solvation shell. Survey of structural data in various crystals containing divalent magnesium ion shows that, in most cases, the coordination number of magnesium Mg$^{2+}$ in crystal structures tends to be six. Ab initio molecular orbital calculations revealed that the lowest energy configuration of hydrated Mg$^{2+}$ has six water molecules packed into the inner shell. For even larger clusters such as MgOCH$_3^+$ (CH$_3$OH)$_6$, the sixth methanol molecule may start to fill the second shell and be hydrogen bonded to the methanol molecules in the first shell.

It has been shown above that the structures for MgOCH$_3^+$ (CH$_3$OH)$_{n-1}$ are significantly different from those of Mg$^{n+}$ (CH$_3$OH)$_n$. While the methanol molecules in the latter tend to be located on one side of Mg$^{2+}$, those in the former series can be evenly distributed around Mg$^{2+}$. The interactions between methanol molecules and Mg$^{2+}$ in MgOCH$_3^+$ (CH$_3$OH)$_n$ are much stronger than those in Mg$^{n+}$ (CH$_3$OH)$_n$, due to the highly polarized MgOCH$_3^+$ molecular ion core. The charge on Mg$^{2+}$ in MgOCH$_3^+$ (CH$_3$OH)$_n$ (1 ≤ n ≤ 5) is between 1.887 and 1.828 and that on the oxygen atom is about −1.25. This contrasts sharply with Mg$^{n+}$ (CH$_3$OH)$_n$, where the charges on Mg and the oxygen atoms are in the range of 0.98–0.95 and of −0.98 to −0.94, respectively.

3. H-Elimination Energies for Mg$^{n+}$ (CH$_3$OH)$_n$ (1 ≤ n ≤ 5). The calculated energetic information on cluster ions

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**TABLE 3: Calculated Energies for Different Structure for MgOCH$_3^+$ (CH$_3$OH)$_n$ ($n$ ≤ 5) at the SCF/6-31G* Level as Well as the Relative Energies for Different Isomers**

<table>
<thead>
<tr>
<th>cluster</th>
<th>structure</th>
<th>relative energies (kJ/mol)</th>
<th>ZPE</th>
</tr>
</thead>
<tbody>
<tr>
<td>MgOCH$_3^+$</td>
<td>I</td>
<td>−313.817 299 419</td>
<td>0.044 785</td>
</tr>
<tr>
<td>MgOCH$_3^+$ (CH$_3$OH)</td>
<td>II</td>
<td>−428.949 218 396</td>
<td>0.102 307</td>
</tr>
<tr>
<td>MgOCH$_3^+$ (CH$_3$OH)$_n$</td>
<td>III</td>
<td>−544.053 800 392</td>
<td>0.160 118</td>
</tr>
<tr>
<td>MgOCH$_3^+$ (CH$_3$OH)$_4$</td>
<td>IV</td>
<td>−659.143 288 599</td>
<td>0.217 493</td>
</tr>
<tr>
<td>MgOCH$_3^+$ (CH$_3$OH)$_6$</td>
<td>Va</td>
<td>−774.212 217 150</td>
<td>0.276 108</td>
</tr>
<tr>
<td>MgOCH$_3^+$ (CH$_3$OH)$_6$</td>
<td>Vb</td>
<td>−774.207 834 400</td>
<td>11.52</td>
</tr>
</tbody>
</table>

a ZPE represents the zero-point energy corrections to the total energies for the most stable isomers.

for MgOCH$_3^+$ (CH$_3$OH)$_n$. In structure II, the O–Mg–O angle is close to 180°. Since Mg$^{2+}$ does not have the single 3s electron as in Mg$^{+}$ (CH$_3$OH)$_n$, the ligand can be evenly located around the central Mg$^{2+}$. Another feature worthy of note is the smaller (CH$_3$OMg)$^+$–CH$_3$OH bond length compared with that in Mg$^{+}$ (CH$_3$OH)$_n$. Clearly, this results from the much stronger interaction between Mg and oxygen due to the highly polarized MgOCH$_3^+$ core. The interaction between Mg and CH$_3$OH can still be considered as electrostatic, but is much stronger here due to the double charge of Mg$^{2+}$. This also applies to larger clusters. In structure III for MgOCH$_3^+$ (CH$_3$OH)$_n$, the calculation shows that the distance between Mg$^{2+}$ and the oxygen atom in CH$_3$OH (1.995 Å) is smaller than that in Mg$^{+}$ (CH$_3$OH)$_n$. Here the three oxygen atoms are almost evenly located around Mg within the same plane. The angles of CH$_3$OH–Mg–OCH$_3$ and CH$_3$OH–Mg–OHCH$_3$ are close to 120°. The optimized structure VI for MgOCH$_3^+$ (CH$_3$OH)$_n$ has the three methanol molecules directly bonded to MgOCH$_3^+$.
TABLE 4. Calculated H-Elimination Energies for Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} (n = 1–5) at the SCF/6-31G* Level\textsuperscript{a}

<table>
<thead>
<tr>
<th>reactions</th>
<th>(\Delta E_0) kcal/mol</th>
<th>(\Delta E_0) eV</th>
<th>(\Delta E_{ZPV}) kcal/mol</th>
<th>(\Delta E_{ZPV}) eV</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mg\textsuperscript{+}(CH\textsubscript{3}OH) \rightarrow MgOCH\textsubscript{3} \textsuperscript{+} + H</td>
<td>82.432</td>
<td>3.575</td>
<td>74.644</td>
<td>3.237</td>
</tr>
<tr>
<td>Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{2} \rightarrow MgOCH\textsubscript{3} \textsuperscript{+} + (CH\textsubscript{3}OH) + H</td>
<td>52.009</td>
<td>2.256</td>
<td>44.010</td>
<td>1.908</td>
</tr>
<tr>
<td>Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{3} \rightarrow MgOCH\textsubscript{3} \textsuperscript{+} + (CH\textsubscript{3}OH)\textsubscript{2} + H</td>
<td>31.015</td>
<td>1.345</td>
<td>23.326</td>
<td>1.011</td>
</tr>
<tr>
<td>Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{4} \rightarrow MgOCH\textsubscript{3} \textsuperscript{+} + (CH\textsubscript{3}OH)\textsubscript{3} + H</td>
<td>16.078</td>
<td>0.697</td>
<td>7.392</td>
<td>0.321</td>
</tr>
<tr>
<td>Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{5} \rightarrow MgOCH\textsubscript{3} \textsuperscript{+} + (CH\textsubscript{3}OH)\textsubscript{4} + H</td>
<td>10.05</td>
<td>0.436</td>
<td>1.258</td>
<td>0.055</td>
</tr>
</tbody>
</table>

\textsuperscript{a} \(\Delta E_0\) and \(\Delta E_{ZPV}\) represent the energies without and with zero-point vibration energy corrections, respectively.

\(\Delta E_0\) and \(\Delta E_{ZPV}\) represent the energies without and with zero-point vibration energy corrections, respectively.

VI. Discussion

1. General Considerations. A potential complication for the laser-ablation source is that the laser-ablated metal ions may not necessarily be in their ground electronic states. The first excited state of the alkaline earth metal ions stems from the promotion of the valence electron to the p- or d-orbitals. It turns out that these np excited states of the metal ions all have a very short lifetime. For example, the lifetime for Mg\textsuperscript{+}(2P) is only 3.7 ns.\textsuperscript{35} Thus, the contribution from these excited states is minimal. Other excited states (e.g., 2D) may participate in the H-elimination reactions. While we cannot exclude any possible reaction of these electronically excited state metal ions, we believe that the observed reaction patterns are mainly from the ground-state reaction based on the following considerations.

The metal ions in the excited states should display different reaction behavior compared with that in the ground states. The ground-state reaction of Mg\textsuperscript{+}(2S) with CH\textsubscript{3}OH to form MgOH\textsuperscript{+} and CH\textsubscript{3} is endothermic by 16 kcal/mol.\textsuperscript{1} Such an endothermic reaction is not expected to occur for the ground-state magnesium ion. However, the ions at higher excited states are energetic enough to react with CH\textsubscript{3}OH to produce MgOH\textsuperscript{+} as the lowest excited-state Mg\textsuperscript{+}(2P) is 102.2 kcal/mol higher in energy than Mg\textsuperscript{+}(2S).\textsuperscript{23} Our experiment did not identify any signals coming from MgOH\textsuperscript{+}. The fact that we only observed the association product Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{5} argues against the possibility of the excited-state reactions in our experiments. In addition, for the dehydrogenation reaction of Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} if the excited-state species (Mg\textsuperscript{+})\textsuperscript{n} were present, the H-elimination reactions would have started at a smaller cluster size. The energy difference between the ground-state Mg\textsuperscript{+}(CH\textsubscript{3}OH) and MgOCH\textsubscript{3} \textsuperscript{+} + H is calculated to be 3.237 eV (74.6 kcal/mol) at the SCF/6-31G* level. Since the three lowest excited states of Mg\textsuperscript{+}(CH\textsubscript{3}OH) are 79.4, 83.9, and 114.5 kcal/mol higher in energy than the ground-state Mg\textsuperscript{+}(CH\textsubscript{3}OH),\textsuperscript{9} the dehydrogenation reaction for Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} may take place for \(n = 1\) or 2 with the electronically excited metal ions. However, no significant amount of H-elimination products MgOCH\textsubscript{3} \textsuperscript{+} + (CH\textsubscript{3}OH)\textsubscript{n} was detected for \(n < 5\), and this lends to the support of our hypothesis that electronically excited Mg\textsuperscript{+} ions did not significantly contribute to the H-elimination reaction.

Another problem with our experiments arises from the kinetic energies of the laser-ablated metal ions as they usually have a wide distribution. While the kinetic energies of metal ions produced from laser ablation are generally in the range 0.1–10 eV,\textsuperscript{36} values as large as 100 eV have been reported depending...
Ion beam experiments have demonstrated that the reactions may become accessible with the argument of ion kinetic energy. The abundant energy of ablated species may affect the reaction kinetics dramatically. In particular, endothermic reactions may be unchanged within the observed kinetic energy range from 10 to 60 eV. As suggested by the authors for the H-elimination reaction. Since H and D are electronically identical, the only differences are associated with the masses and are thence primarily with the zero-point vibrational energies. For Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} and Mg\textsuperscript{+}(CH\textsubscript{3}OD)\textsubscript{n}, the number of OH(OD) bonds in MgOCH\textsubscript{3}\textsuperscript{+}(CH\textsubscript{2}OH)\textsubscript{n-1} (MgOCH\textsubscript{3}\textsuperscript{+}(CD\textsubscript{2}OD)\textsubscript{n-1}) is less than those in Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} (Mg\textsuperscript{+}(CD\textsubscript{2}OD)\textsubscript{n}) by one. The energy drop from MgOCH\textsubscript{3}\textsuperscript{+}(CH\textsubscript{2}OH)\textsubscript{n-1} to MgOCH\textsubscript{3}\textsuperscript{+}(CD\textsubscript{2}OD)\textsubscript{n-1} is smaller than that from Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} to Mg\textsuperscript{+}(CH\textsubscript{2}OD)\textsubscript{n}. As a result, more methanol molecules are needed to reach the switching point from Mg\textsuperscript{+}(CH\textsubscript{2}OD)\textsubscript{n} to MgOCH\textsubscript{3}\textsuperscript{+}(CH\textsubscript{2}OH)\textsubscript{n-1} than those needed for the Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} system. The situation for Mg\textsuperscript{+}(CD\textsubscript{2}OD)\textsubscript{n} is similar to that for Mg\textsuperscript{+}(CH\textsubscript{2}OD)\textsubscript{n}. For the Mg\textsuperscript{+}(CH\textsubscript{2}OD)\textsubscript{n} and Mg\textsuperscript{+}(CD\textsubscript{2}OD)\textsubscript{n} systems, however, there should be no isotope effect for the dehydrogenation reaction since no bond breaking or formation process occurs to the methyl group in the reaction.

In connection to the first product switching, we want to mention the recent work by Stace and co-workers on the gas-phase chemistry of Mg\textsuperscript{+} and Mg\textsuperscript{2+} with methanol clusters with a different experiment setup. Neutral magnesium—methanol clusters were formed through the reaction of the methanol clusters with a magnesium vapor. The clusters were then subjected to 100 eV electron impact ionization. The singly and doubly charged cluster ions Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n} and Mg\textsuperscript{2+}(CH\textsubscript{3}OH)\textsubscript{n} were generated. They also observed the ion signals corresponding to MgOCH\textsubscript{3}\textsuperscript{+}(CH\textsubscript{2}OH)\textsubscript{n} for n ≥ 3 and ascribed them to the result of the dehydrogenation of Mg\textsuperscript{+}(CH\textsubscript{3}OH)\textsubscript{n}. This critical size is then much smaller than what we found in the present work. There are two possible reasons for this discrepancy. First, the electron impact ionization they used is usually considered as a “nonsoft” ionization technique and may therefore result in high internal excitation in the clusters. The high internal energies of [Mg(CH\textsubscript{3}OH)\textsubscript{n}]\textsuperscript{+} may reduce the critical size from n ≥ 5 to n ≥ 3. Second, it is possible that the dehydrogenation products MgOCH\textsubscript{3}\textsuperscript{+}(CH\textsubscript{2}OH)\textsubscript{n-1} in Stace’s experiment are generated from the deprotonation of the doubly charged ions [Mg(CH\textsubscript{3}OH)\textsubscript{n}]\textsuperscript{2+}. Previous studies on doubly charged alkaline earth metal ions—water clusters by Kebarle et al. showed that these cluster ions undergo collision-induced dissociation (CID) and give charge reduction products at low n. For example, the reaction of Mg\textsuperscript{2+}(H\textsubscript{2}O)\textsubscript{n} to give MgOH\textsuperscript{+}(H\textsubscript{2}O)\textsubscript{h} + H\textsubscript{2}O\textsuperscript{+} occurs for n ≥ 3.

One of the most elusive aspects about the H-elimination reactions is their mechanism. On the basis of our experimental results and chemical intuition, we believe that the reaction corresponding to the first switching is through a complex mechanism, which is different from that for the reaction of the methanol monomer with Ca\textsuperscript{+}, Sr\textsuperscript{+}, and Ba\textsuperscript{+}. Initially, the reaction complex is formed and accompanied perhaps by evaporation of solvent molecules. At a certain stage, the electron transfer from the metal center to the methanol molecules must take place, resulting in the H-elimination.
As in the first product switching, we discuss the second product switching based on the assumption that the critical size is determined by thermodynamic factors. Alternatively, one might emphasize the importance of the hydrogen transport or tunneling dynamics in the clusters. For the reaction within such a large methanol cluster as Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\), methanol molecules surrounding the metal ions may play the role of a solvation cage, which would create an energy barrier along the reaction path for H-elimination. Conceivably, the trapping probability of the eliminated H atom upon reaction should increase with increasing cluster size. As a result, the H-elimination reaction may be suppressed for sufficiently large clusters. In fact, these two explanations are inter-related since the stability of the H-trapped structure will certainly affect the H-trapping probability.

The key to understanding the second product switching is the structures of the products before and after the product switching. For the first product switching, it is quite clear that the metal ion–solvent cluster structure has turned to the H-elimination products with the formation of a MOCH\(_3\)\(^+\) ion core. The second switching is more ambiguous in this regard. Since after the second product switching, the products have the same stoichiometry as that of the association products without H-elimination, one would speculate that their structures may also be the same. However, this seems inconsistent with the thermodynamic reasoning as well as our experimental results. First, consider two energy curves as a function of cluster size: one corresponds to the H-elimination products and the other corresponds to the association products. In the mass region below the critical size for the first product switching, the energies of MgOCH\(_3\)\(^+\)·(CH\(_3\)OH)\(_n\) are higher than those of Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\). Since the energy decrease of Mg-OCH\(_3\)\(^+\)·(CH\(_3\)OH)\(_{n-1}\) is faster than that of Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\) with cluster size, the two energy curves will cross, leading to the first product switching. But if the energies of the products are monotonic and continuous functions of the cluster size, one would expect the two curves to meet again at a somewhat larger cluster size, which would constitute the second product switching.

Another important clue concerning the product structures after second product switching comes from the deuterium substitution effect. For clusters below the first product switching cluster size, where MgOCH\(_3\)\(^+\)·(CH\(_3\)OH)\(_n\) is higher than Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\) in energy, replacing OH by OD makes the energy difference even larger due to the zero-point energy corrections. It then requires more methanol solvation for the H-elimination reaction to occur. This is what we observed in our experiments. For the second product switching, our experiments showed that the switching cluster size is smaller upon deuterium substitution (from \(n = 15\) to \(n = 14\)). This suggests that the products created after the second product switching may not involve the H–O bond severance. The reason is that the deuterium substitution will shift the energy curve of MgOCH\(_3\)\(^+\)·(CH\(_3\)OH)\(_n\) upward relative to the energy curve of Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\), and have no effect on the product energy curve if the H–O bond cleavage does not occur. The consequence is that the switching cluster size will be smaller. This fits nicely into the results of our isotope substitution experiments. If, on the other hand, the product structures embody the H–O bond cleavage, the deuterium substitution will shift both product energy curves to a similar extent. The net outcome is that the switching point remains the same and there would be no isotope effect for the second product switching in this case.

The above considerations carry two crucial points: (1) the product structures should not involve H–O bond cleavage, for example, MgOCH\(_3\)\(^+\)·H\(_2\)O\(_n\)·H\(_2\)O\(_n\), a similar structure as that proposed for the Mg\(^{2+}\)·H\(_2\)O system, and (2) the product structures should be different from those of the association products Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\). The burning question is then, what are the product structures on earth after the second product switching?

Remember for the first product switching, the reason Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\) is relatively unstable with increasing cluster size compared with MgOCH\(_3\)\(^+\)·(CH\(_3\)OH)\(_{n-1}\) is that Mg in the former species has a valence electron around it and the oxidation state of Mg is +1, while in the latter case Mg is oxidized to Mg\(^{2+}\). Therefore, the electrostatic interactions between the methanol ligands and the central Mg ion in the latter complex are much stronger than those in the former complex such that the energy needed for transferring the electron from Mg to the methanol ligands is sufficiently compensated at a certain cluster size. For the second product switching, we believe that the electron transfer from the Mg to the methanol ligands did take place as in the first product switching. But in this case, H-elimination does not occur. This is reasonable because the free electron can be stabilized by solvation given sufficient number of methanol molecules. Ultimately, Mg must exist in the form of Mg\(^{2+}\) in the solution of polar solvents. In addition, this is consistent with the previous inferences that the product structures should have all their H–O bonds intact, and they also should be different from those of the association products.

To proceed a step further, we conjecture a two-cage structure for the products beyond the second switching. In this novel structure, one cage is centered on the Mg\(^{2+}\) and the other is around the free electron. For the metal ion Mg\(^{2+}\), since it has an oxidation state of +2 without outer valence electrons, it should be able to have the methanol ligands evenly distributed around Mg\(^{2+}\). The number of methanol ligands in the first solvation shell is expected to be six based on the knowledge on the coordination of Mg\(^{2+}\) in crystal. The two solvation cages can be connected by intermediate methanol molecules through hydrogen bonds. Clearly, such a structure should be unstable at small cluster sizes because there is not enough methanol molecules to solvate the transferred electron. Only when the cluster reaches a certain size can this structure then compete favorably with that of the H-elimination products. It should be pointed out that for the solvated electron cage, the electron may not necessarily be in the center. Much further work is needed to understand the structure in greater detail.

The two-cage structure is similar to the previously proposed ion-pair structure of Sr\(^{2+}\)·(NH\(_3\))\(_n\) by Fuke et al.48,49 Also proposed such an ion-pair structure for the Mg\(^{2+}\)·(H\(_2\)O)\(_n\) system as one of the candidates to explain the second switching behavior. However, the detailed picture for the ion-pair structure has not yet been worked out. The two-cage structure we proposed should be correctly written as [Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\)]\(e^-\)·(CH\(_3\)OH)\(_n\) rather than the simple Mg\(^{2+}\)·(CH\(_3\)OH)\(_n\). Experimentally, there has been no report on the negatively charged methanol clusters in the gas phase. However, for water and ammonia, stable negatively charged cluster ions were observed when the clusters reached a critical size.50,51 It is reasonable to expect that such a kind of methanol cluster anions also exists if enough methanol...
molecules are available to solvate the electron, in much the same way as the solvated electrons in low-temperature methanol glass.\(^{45}\) Moreover, the two-cage structure is further stabilized by the attractive Coulombic interaction between the two cages.

The electrostatic forces between \(\text{Mg}^{2+}\) and polar ligands such as water and methanol correspond to the charge-dipole interactions, which scale as \((Zalr^2)\). The larger the dipole moment of the ligand, the larger the interactions. For the solvated electron, the interactions between the electron and solvent molecules are also of a charge–dipole type.\(^{45,52}\) Under the same condition, the complex with solvent molecules of larger dipole moments will be more stable than those with smaller dipole moments. It is known that the dipole moment of methanol is 1.7 D and that of water is 1.85 D.\(^{23}\) Qualitatively, more methanol molecules may be required to stabilize the two-cage structure for the metal ion–methanol clusters than the number of water molecules, which are needed for the metal ion–water clusters. Experimentally, we found the critical size for the second product switching of the metal-ion-methanol cluster system to be \(n = 15\) for both \(\text{Mg}\) and \(\text{Ca}\), which is larger than that of the metal ion–water cluster system (the switching occurs at \(n = 13\) for \(\text{Ca}\) and \(n = 15\) for \(\text{Mg}\)). This observation seems to be fairly consistent with the charge-dipole nature for the interactions, which is assumed to be operative in the metal ion–solvent cluster systems.

4. Comparison with the \(\text{M}^{n+}-(\text{H}_2\text{O})_n\) System (\(\text{M} = \text{Mg}\) and \(\text{Ca}\)). The reaction system \(\text{M}^{n+}-(\text{CH}_3\text{OH})_n\) in the present study is remarkably similar to the \(\text{M}^{n+}-(\text{H}_2\text{O})_n\) system in the sense that both the first and the second switchings take place in the same cluster size regions. This is surprising considering that, although \(\text{CH}_3\text{OH}\) and \(\text{H}_2\text{O}\) are both polar molecules, their solvent cluster structures are expected to be quite different. \(\text{H}_2\text{O}\) has two identical H-atoms and both can be used for the hydrogen-bonded networks in water clusters, whereas only one H-atom in \(\text{CH}_3\text{OH}\) can be used for hydrogen bonding. One should note that the structures of the pure solvent clusters are very different from those in \(\text{M}^{n+}-(\text{solvent})_n\) and the solvated H-elimination products. It is the structures of \(\text{M}^{n+}-(\text{solvent})_n\) and the solvated H-elimination products, instead of the pure solvent clusters, that are important in dictating the H-elimination energies. This suggests a plausible explanation for the similarity in the H-elimination patterns between \(\text{M}^{2+}-(\text{CH}_3\text{OH})_n\) and \(\text{M}^{+}-(\text{H}_2\text{O})_n\). This explanation assumes that, in the cluster range of our investigation, only one H–O bond of \(\text{H}_2\text{O}\) is actively involved in the H-bonding in both \(\text{M}^{+}-(\text{H}_2\text{O})_n\) and the solvated H-elimination products. The other H–O bond, being idle, can be replaced by \(\text{CH}_3\)–O without much effect on the cluster structure. This is indeed what we found in our ab initio calculations. The structures of \(\text{M}^{+}-(\text{CH}_3\text{OH})_n\) and \(\text{M}^{+}-(\text{H}_2\text{O})_n\) are quite similar. They both have three molecules in the first solvation shell; the second shell is attached to the first shell through hydrogen bonding. The structures of the H-elimination products are also similar for the two metal ion–solvent cluster systems.

VII. Summary

We have presented the results of our studies on the gas-phase reactions between singly charged alkaline earth metal ions (\(\text{Mg}^{+}\), \(\text{Ca}^{+}\), \(\text{Sr}^{+}\), and \(\text{Ba}^{+}\)) and methanol clusters in a pick-up cluster source. The reaction products are shown to exhibit anomalous size distributions characterized by two product switching cluster size regions, analogous to the reaction behaviors of the alkaline earth metal ion–water cluster systems. The first product switching from \(\text{M}^{+}(\text{CH}_3\text{OH})_n\) to \(\text{MOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) occurs for \(n\) at around 5. The \(\text{MOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) species switches back to \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\) for \(n\) at around 15 (second switching). The critical sizes for the first and the second switching were found to be affected by the metal and deuterium substitutions. Deuterium substitution on the OH group is shown to affect the critical sizes for both product switchings, which is explained based on the thermodynamics of the H-elimination reactions.

The reason for the first product switching is ascribed to the different thermodynamic stabilities for the two relevant species with increasing cluster size. In \(\text{MOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) cluster ions, Mg is oxidized to \(\text{Mg}^{2+}\) and the strong electrostatic interactions between \(\text{Mg}^{2+}\) and the methanol ligands strongly stabilize the system. This is in contrast to \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\), where the oxidation state of Mg is +1. The \(\text{MgOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) system is stabilized much more quickly than \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\) with the increasing cluster size, and the transition between the two species readily occurs when the cluster reaches the critical size. For the second switching from \(\text{MOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) to \(\text{M}^{+}(\text{CH}_3\text{OH})_n\) at large cluster sizes, we believe that the later series has a structure different from those of the small cluster ions. These large clusters are assumed to have an ion-pair structure. In particular, we have proposed a two-cage structure \([\text{Mg}^{2+}(\text{CH}_3\text{OH})_n][\text{e}^{-}(\text{CH}_3\text{OH})_m]\), which embodies both ion solvation and electron solvation.

Structures of singly charged magnesium ion-methanol clusters \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\) and their H-elimination products \(\text{MgOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) have been studied by ab initio calculations at the SCF/6-31G* level. The association product series and the H-elimination product series are found to have dramatically different structures. For \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\), the methanol ligands are located on one side of Mg to avoid the high electron density on the other side resulting from the single valence electron of \(\text{Mg}^{+}\). The number of methanol molecules that can be directly bonded to \(\text{Mg}^{+}\) is affected by the ligand–ligand repulsions and is limited to three. They form the first solvation shell and the fourth methanol will start to fill the second shell. For \(\text{MgOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\), the methanol molecules can be evenly located around Mg and the natural charge calculation found the charge of Mg to be close to +2. As a result, the electrostatic interactions between Mg and methanol ligands are much stronger than those in \(\text{Mg}^+(\text{CH}_3\text{OH})_n\), where the Mg has an oxidation state of +1. Consequently, the \(\text{MgOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) series is stabilized much more quickly than \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\), with the increasing cluster size. The energy change for the H-elimination reaction of \(\text{Mg}^+(\text{CH}_3\text{OH})_n\) decreases from 3.237 eV for \(n = 1\) to 0.055 eV for \(n = 5\). Qualitatively, this is consistent with our experimental results that \(\text{Mg}^{+}(\text{CH}_3\text{OH})_n\) starts to change to \(\text{MgOCH}_3^{+}(\text{CH}_3\text{OH})_{n-1}\) for \(n = 5\). The driving force for the H-elimination reaction appears to be related to the fact that the products are thermodynamically more stable than the reactants at large cluster sizes.

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References and Notes
