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Bonded Network**

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## ARTICLE

# Investigating the Hydrolysis of Cryogenically Layered Molybdenum Hexafluoride Through a Disordered Hydrogen-Bonded Network

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Molybdenum hexafluoride (MoF<sub>6</sub>) is used as a non-radioactive substitute for uranium to study the hydrolysis of metal hexafluorides. Molybdenum hexafluoride gas and water vapor, from the air, were sequentially layered onto a diamond substrate kept at liquid nitrogen temperature using a custom designed cryogenic cell with a copper cold finger. Reaction progress was monitored by transmission Fourier Transform Infrared Spectroscopy (FTIR) through the layers and diamond substrate over several hours while allowing the substrate to warm. Changes in the modes in the 500–1000 cm<sup>-1</sup> region are tracked as the reaction progresses in order to identify intermediate species. Strong absorption features are also observed in the 1000–3000 cm<sup>-1</sup> range, suggesting the presence of ionic dissociation intermediates trapped in a disordered H-bonded network of cryogenic hydrofluoric acid. A possible reaction pathway is proposed and the final hydrolysis product is characterized by FTIR, UV-Vis, and Scanning Electron Microscopy/Energy Dispersive X-Ray Spectroscopy (SEM/EDS).

## Introduction

Uranium hexafluoride (UF<sub>6</sub>) is a highly utilized material feedstock for uranium enrichment processes, including gas diffusion, gas centrifuge, and some laser separations. These enrichment processes take advantage of the solid material state of UF<sub>6</sub> at room temperatures and ambient pressures, as well as the ease of sublimation upon heating. As a result of the common use of UF<sub>6</sub> in uranium enrichment since the 1940s, the chemical processing of UF<sub>6</sub> has been highly studied and is well understood<sup>1-8</sup>. However, uranium enrichment is carried-out under controlled conditions to limit moisture in the process to prevent loss of uranium to more thermodynamically favorable reactions which produce solids (material holdup). When UF<sub>6</sub> is exposed to atmosphere, it readily undergoes the hydrolysis reaction  $\text{UF}_6(\text{g}) + 2\text{H}_2\text{O}(\text{g}) \rightarrow \text{UO}_2\text{F}_2(\text{s}) + 4\text{HF}(\text{g})$ . The reaction pathway of the UF<sub>6</sub> hydrolysis reaction is not well understood, and computationally predicted key intermediates have never been observed experimentally.

Previous research performed at Oak Ridge National Laboratory on uranyl fluoride (UO<sub>2</sub>F<sub>2</sub>) particulate formation suggests the final product of the reaction is a function of the relative humidity, where small fractal aggregates form when a limited amount of water is available and larger spheroidal

particles form at high relative humidity<sup>9, 10</sup>. Historically, research has focused on gas phase spectroscopy to better understand the chemical products of the reaction<sup>11-13</sup>, although several studies have characterized the chemical morphology and speciation of the products<sup>14-17</sup>. More significant effort has been applied to computational studies which have aimed to elucidate the reaction dynamics of the hydrolysis reaction<sup>18-24</sup>. Computational studies have primarily predicted the hydrolysis reaction progression through experimentally unobserved intermediates, most commonly UF<sub>5</sub>OH.

There is a very limited amount of research on whether the hydrolysis reaction is required to occur: (1) in the gas-phase, creating UO<sub>2</sub>F<sub>2</sub> molecules which then cluster to form larger particulates, (2) if initial formation of molecular UO<sub>2</sub>F<sub>2</sub> can seed the formation of clusters at earlier times and the reaction then proceeds freely in the condensed phase, or (3) if both pathways are possible. With limited experimental data on the hydrolysis reaction in the condensed phase and no thermodynamic data for UF<sub>5</sub>OH, computational studies are unable to definitively assess the role of condensed phase chemistry<sup>24</sup>.

In the work described here, the hydrolysis of molybdenum hexafluoride (MoF<sub>6</sub>) was studied as a surrogate reaction for the hydrolysis of UF<sub>6</sub> to reduce safety concerns regarding the handling of radioactive material. MoF<sub>6</sub> also readily undergoes hydrolysis when in contact with atmospheric humidity and is a known contaminate in nuclear fuel cycle byproducts. The hydrolysis reaction was studied in the condensed phase under cryogenic conditions to slow down the reaction kinetics and capture possible reaction intermediates. Since particulate formation has only been studied in the gas phase, working in the condensed phase will also elucidate what role condensed phase reactions may have on product particulate formation. A novel

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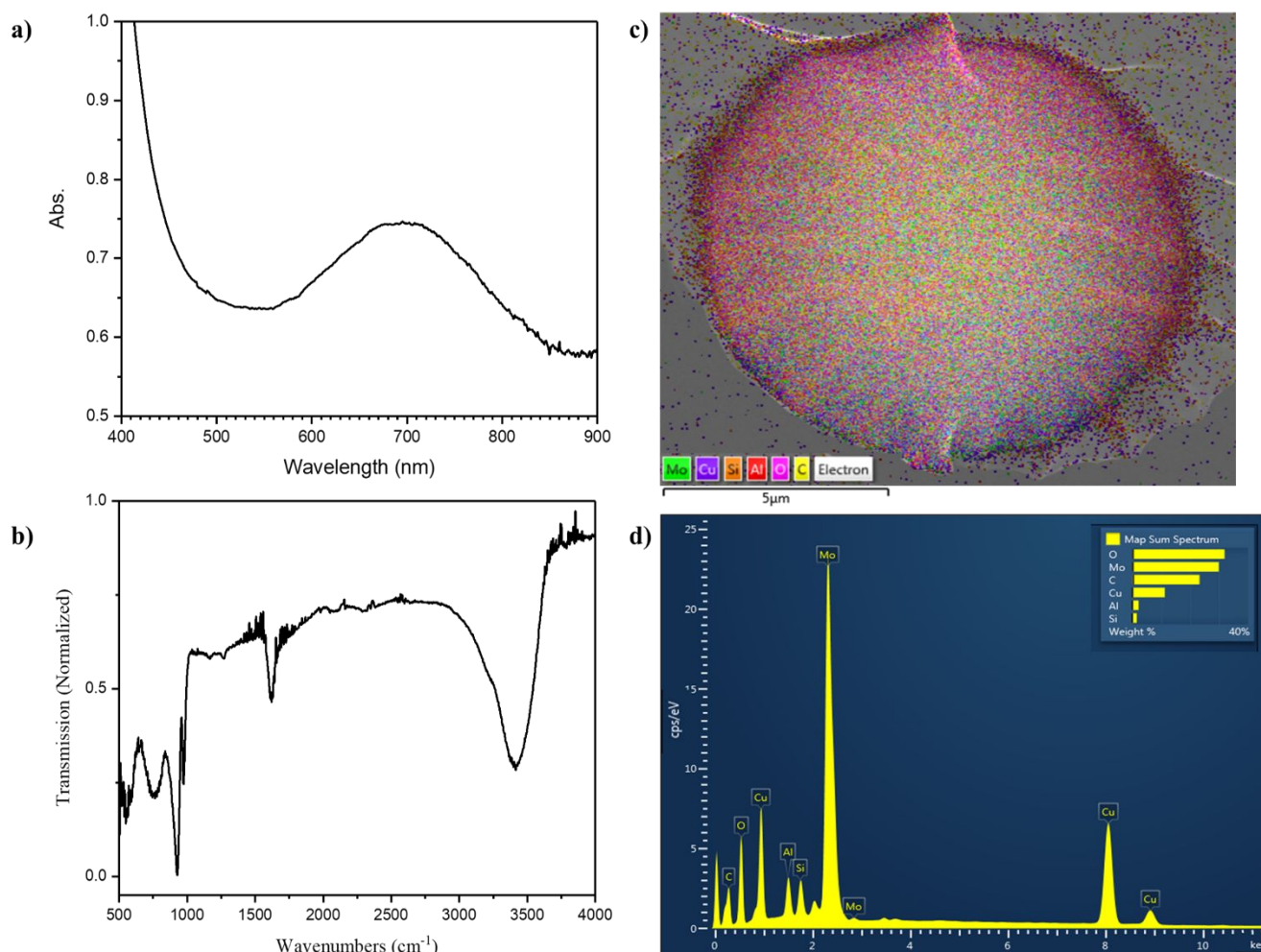


Figure 1 a) UV/visible absorption spectrum of final product dissolved in DI water, b) Transmission FTIR spectrum of product on diamond substrate, c) EDS layered image of product, d) EDS graph of product (Cu, C, Si, and Al are all artefacts from the stage and stage holder).

cryogenic layering technique was used to freeze the reactants to prevent spontaneous hydrolysis and control the rate of the reaction.

The hydrolysis of  $\text{MoF}_6$  has received little attention in the literature and this work intends to: (1) demonstrate the ability of our layering technique to track and isolate reaction intermediates, (2) elucidate the reaction pathway governing the hydrolysis of  $\text{MoF}_6$ , and (3) use the information learned about the hydrolysis of  $\text{MoF}_6$  to better understand possible reaction pathways of  $\text{UF}_6$  hydrolysis.

## Results and Discussion

### $\text{MoF}_6$ Hydrolysis Product Characterization

The final product of the hydrolysis of  $\text{MoF}_6$  in the condensed phase cryogenic layering experiment was a deep blue film covering the diamond substrate. If  $\text{MoF}_6$  hydrolysis were a perfect substitute for  $\text{UF}_6$ , the expected final product would be  $\text{MoO}_2\text{F}_2$ . However, the product was identified as a hydrate of  $\text{MoO}_3$  based on the characteristics of previously reported molybdenum oxide samples in the literature<sup>25-33</sup>. In addition to

FTIR, the product was also characterized using UV-Vis and SEM/EDS (Figure 1).

### $\text{MoF}_6$ Hydrolysis by Cryogenic Layering

After sequential layering of  $\text{MoF}_6$  and air, the spectrum (figure 2) initially displays the expected absorption bands associated with condensed phase water and  $\text{CO}_2$ : coupled intramolecular OH stretching modes around  $3250\text{ cm}^{-1}$ , HOH bending modes at  $1700\text{ cm}^{-1}$ , intermolecular librations at  $900\text{ cm}^{-1}$ , the CO asymmetric stretching mode at  $2350\text{ cm}^{-1}$ , and the OCO bending modes at  $550\text{ cm}^{-1}$ . In addition, extra bands can be observed in the  $700\text{ cm}^{-1}$  region which can be attributed to the cryogenically deposited  $\text{MoF}_6$ . The deposited  $\text{MoF}_6$  exhibits three peaks red shifted, but spectrally similar, to gas phase  $\text{MoF}_6$ . Based on previous work<sup>34-36</sup>, these peaks are identified as the  $\nu_3$  fundamental at  $700\text{ cm}^{-1}$ , the  $2\nu_4 + \nu_6$  combination band at  $715\text{ cm}^{-1}$ , and the  $3\nu_4$  overtone as a shoulder at  $770\text{ cm}^{-1}$ .

Once the diamond substrate begins to warm, significant changes in the spectral features of all regions are observed. Initial changes in the  $\text{MoF}_6$  spectral region are difficult to discern from simple phase changes due to the increase in temperature. Similar spectral changes were observed in the hydrolysis

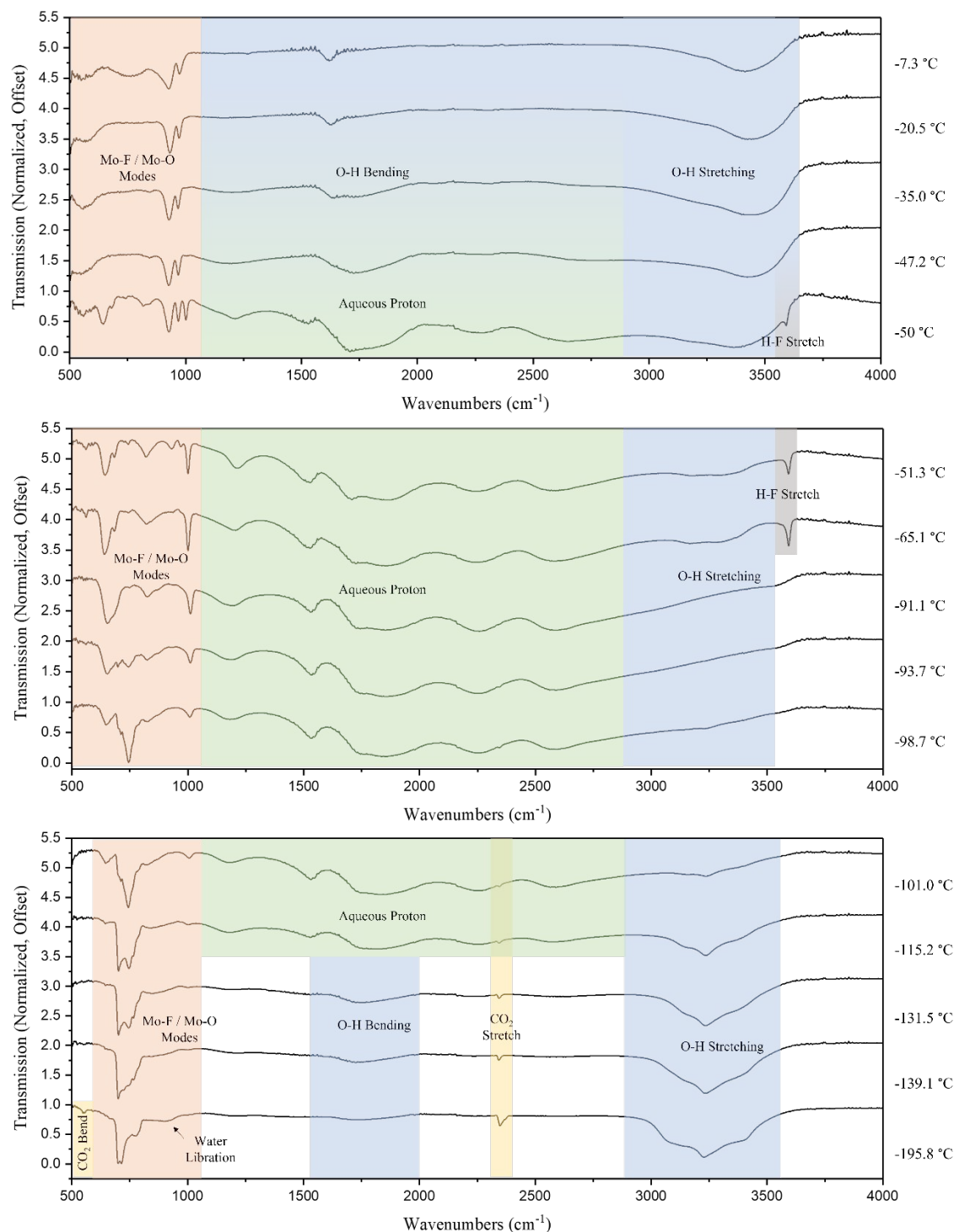


Figure 2 FTIR spectrum of the  $\text{MoF}_6$  hydrolysis as temperature rises. Different regions have been color coded to correspond to known bands of water (blue),  $\text{CO}_2$  (yellow), aqueous proton (green), and molybdenum fluorides/oxides (red). The experiment starts at  $-197.8$  °C and warms up to  $-7.3$  °C over the course of 2 hours.

reaction experiment and as changes observed upon allowing cryogenically deposited  $\text{MoF}_6$  to warm up in the absence of water (Figure S1). There is also overlap between the spectral region associated with the intermolecular librations of water and the  $\text{MoF}_6$  modes. These factors make it difficult to identify new species produced during the early steps of the hydrolysis reaction, i.e. at temperatures below  $-100$  °C. However, as the

reaction proceeds, a clear deviation from the pure  $\text{MoF}_6$  warming spectra are observed and reaction intermediates can be identified. As expected, the signature peaks of  $\text{CO}_2$  disappear as the substrate warms up - the  $\text{CO}_2$  sublimates and is pulled off by the vacuum. The  $\text{CO}_2$  sublimation happens concurrently with the depletion of the water OH stretching modes past  $3000$   $\text{cm}^{-1}$  and the in-growth of a broad continuous absorption band from

roughly 1000-3000  $\text{cm}^{-1}$ . The broad continuum of peaks in this intermediate region persists through the reaction until the final product is formed.

The continuum observed during the progression of the reaction is the spectroscopic signature of aqueous protons and is spectrally similar to previously reported spectra of HF dissociated in amorphous solid water<sup>37</sup>. Very broad and intense absorbance features initially appear around 1200  $\text{cm}^{-1}$ , 1500  $\text{cm}^{-1}$ , 1850  $\text{cm}^{-1}$ , 2250  $\text{cm}^{-1}$ , and 2550  $\text{cm}^{-1}$ . After more time, two new features grow in around 3300  $\text{cm}^{-1}$  and 3600  $\text{cm}^{-1}$ . The large continuum provides evidence that fluorine atoms have been hydrolyzed from  $\text{MoF}_6$  and are dissolved in an extended hydrogen bonded network. The spectral features of the aqueous proton-type continuum have been heavily studied, though there is still a healthy debate in the literature over the origin of many of these features. The unique method by which the network was created, and reaction monitored, allows for the results to provide some clarity on the debate.

The 1200  $\text{cm}^{-1}$  band lies in an ambiguous spectral range based on previous experimental and computational studies<sup>38-47</sup>. Experimental data and electronic structure calculations suggest the intramolecular acidic proton symmetric stretching and bending modes of proton shared species, as well as several asymmetric vibrational modes of proton shared species and the symmetric bending umbrella mode of the Eigen-type cations are all expected to contribute oscillator strength around 1200  $\text{cm}^{-1}$ . In the observed hydrolysis reaction of  $\text{MoF}_6$ , spectral shifting of the bands position and peak shape are associated with the presence of HF stretching in the extended network. The band blue shifts from 1175  $\text{cm}^{-1}$  to 1215  $\text{cm}^{-1}$  in the presence of HF and becomes noticeably more defined. The band then red shifts back to 1175  $\text{cm}^{-1}$  as the HF is pulled off by vacuum. The changes in the band location suggests a proposed proton shared species such as  $[\text{H}_2\text{O}\cdot\text{HF}](\text{aq})$  are primarily responsible for contributing to the oscillator strength around 1215  $\text{cm}^{-1}$  when an increased concentration of HF appears to be present. Species not directly sharing a proton with HF are then the major contributors around 1175  $\text{cm}^{-1}$ .

The large band centred near 1850  $\text{cm}^{-1}$  overlaps with the OH bending mode around 1700  $\text{cm}^{-1}$  and has historically been assigned to Eigen-type ions asymmetric bending vibrations<sup>39, 41-43, 45, 48</sup>. However, there is literature which also supports the role of acidic proton asymmetric stretching vibrations of proton-shared species as more compelling candidates<sup>37, 47, 49, 50</sup>. Here, the 1850  $\text{cm}^{-1}$  band appears well before the water OH stretching region completely depletes and remains constant through the change in HF concentration, even persisting after the HF has been removed from the reaction. The intensity of this band appears to be independent of HF concentration in the performed experiment and only depends on the presence of fluoride ions, which supports the Giguere and Turrell  $[\text{F}\cdot\text{H}_3\text{O}^+](\text{aq})$  contact ion pair model<sup>51</sup>.

The broad band at 2550  $\text{cm}^{-1}$  is attributed to either product-like Eigen-type  $\text{H}_3\text{O}^+(\text{aq})$  ions, which display a similar spectral feature as observed in amorphous mixtures of the strong acids HCl and HBr<sup>52, 53</sup>, or OH stretching vibrations from strongly distorted water molecules in the first shell of fluoride anions<sup>54-</sup>

<sup>56</sup>. The band has also been attributed to HF molecules bound to tetra-coordinated water<sup>57</sup>. Again, here the band is independent of HF concentration, leading to the conclusion that molecular HF is not directly involved in species contributing to oscillator strength in this area.

The region between 500  $\text{cm}^{-1}$  and 1000  $\text{cm}^{-1}$  is of much greater interest to the authors as it contains the relevant Mo-F and Mo-O stretching modes. Unfortunately, most spectroscopic data for  $\text{MoF}_6$  has been collected in the gas phase, and there is limited research on  $\text{MoF}_6$  hydrolysis, significantly complicating the identification of unique species in this range. The initial peaks around 700-780  $\text{cm}^{-1}$  can be attributed to pure  $\text{MoF}_6$  and identification of the product reveals the peaks in 500-1000  $\text{cm}^{-1}$  region to be associated with  $\text{MoO}_3$  hydrates, providing an overall reaction scheme of  $\text{MoF}_6 + 3\text{H}_2\text{O} \rightarrow \text{MoO}_3 + 6\text{HF}$ . Since the initial steps of the reaction are convoluted by changes in phase, working in reverse simplifies band determinations.

The product shows four distinct bands between 500-1000  $\text{cm}^{-1}$  and two bands associated with water bending and stretching at 1700  $\text{cm}^{-1}$  and 3400  $\text{cm}^{-1}$ , respectively. The two bands between 900-1000  $\text{cm}^{-1}$  can be identified as Mo=O stretching based on the product identification and previous experimental data<sup>25-32</sup>. These two bands appear early in the reaction and can be seen after the presence of HF is detected. As the reaction proceeds, these two peaks grow in stronger and remain the only peaks in the spectral region after HF has been fully removed. The band around 550  $\text{cm}^{-1}$ , attributed to O-Mo-O bending<sup>25-32</sup>, appears much earlier in the reaction and is correlated with the disappearance of HF in the spectra. These three peaks seen in the penultimate stage of the reaction are attributed to the formation of the molecular product  $\text{MoO}_3$  during the final stages of the reaction. At this point in the reaction, the aqueous proton network is depleting due to the evaporation of  $\text{H}_2\text{O}$  and HF, and the  $\text{MoF}_6$  has been fully hydrolyzed to  $\text{MoO}_3$ . The remaining peak at 750  $\text{cm}^{-1}$  grows in much later than the other bands and is typically observed in large aggregates or nanoparticles of  $\text{MoO}_3$  and water. The presence and timing of the 750  $\text{cm}^{-1}$  suggest the band is due to the aggregation of the molecular  $\text{MoO}_3$  hydrate and the formation of particulates in the condensed phase<sup>58-63</sup>.

Moving back a step in the reaction process from the end-product, three distinct bands are present across the intermediate reaction (from approx. -50°C to -100°C), 650  $\text{cm}^{-1}$ , 850  $\text{cm}^{-1}$ , 1000  $\text{cm}^{-1}$ , which all disappear along with the depletion of HF and appear with the formation of the extended aqueous proton network. Based on literature<sup>25, 64-68</sup>, the peak at 1000  $\text{cm}^{-1}$  is consistent with Mo=O stretching in neutral molybdenum fluorido-oxido compounds as well as molybdenum oxide stretching in similar anionic systems. Since our reaction is taking place surrounded by water the intermediates formed will be stabilized by the extended hydrogen bonding network of the aqueous proton. The molybdenum oxygen modes in both neutral and anionic systems are represented here as  $[\text{MoO}/\text{MoO}][\text{H}_{\text{net}}]$ . Since the 650  $\text{cm}^{-1}$  and 850  $\text{cm}^{-1}$  bands are heavily correlated to the mode at 1000  $\text{cm}^{-1}$ , these major bands may be attributed to Mo-F stretching, bridged Mo-O-Mo, and bridged Mo-F-Mo modes.

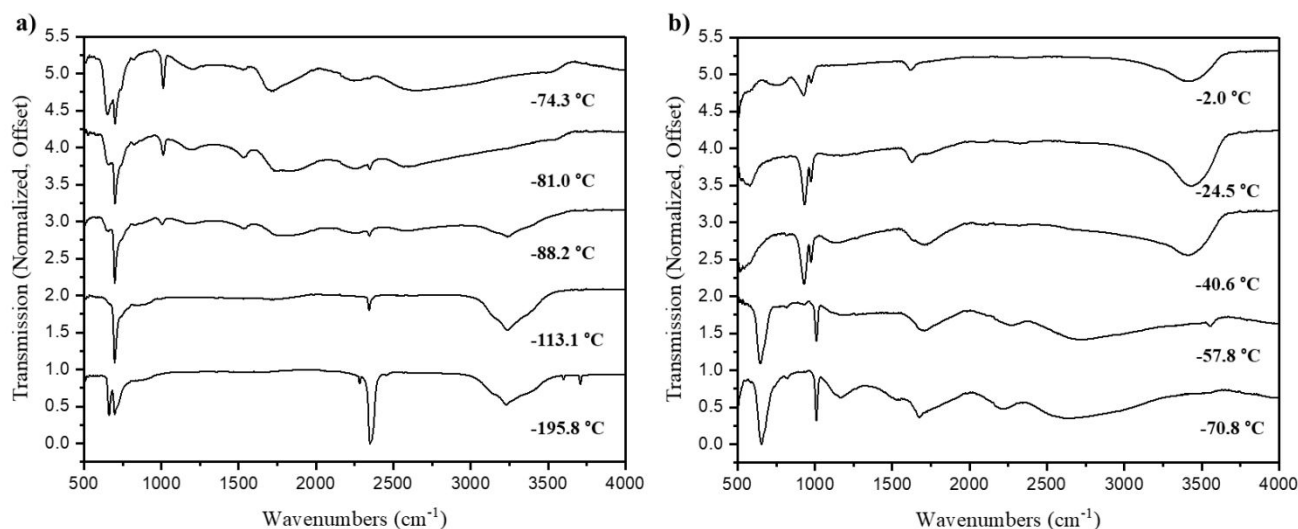


Figure 3 FTIR spectrum of the  $\text{MoF}_6$  hydrolysis once an insulating layer of  $\text{CO}_2$  has been used to initially separate the  $\text{MoF}_6$ . a) spectra of initial unreacted species is followed by depletion of the insulating  $\text{CO}_2$  later and formation of aqueous proton network and b) Reaction proceeds to  $\text{MoO}_3$  product as temperature continues to increase. Temperature is increasing from bottom to top.

Bridged Mo-F-Mo formation is common in these types of copounds; however, the reaction is taking place in a water network, so bridged Mo-O-Mo intermediates are also considered here and may be difficult to spectroscopically differentiate from Mo-F-Mo bridging. There are several other minor peaks and shoulders between  $650\text{--}850\text{ cm}^{-1}$ , which are difficult to characterize due to low intensity and limited literature on the reaction, but are likely associated with intermediate and bridged compounds. Assuming a similar reaction pathway as proposed for  $\text{UF}_6$  hydrolysis, the initial intermediate from the  $\text{MoF}_6$  hydrolysis will likely be  $\text{MoF}_5\text{OH}$ , which should then quickly react to form  $\text{MoOF}_4$ . Knowing the final product to be fully hydrolyzed, fluorine atoms must be continually hydrolyzed, and a range of possible intermediates can be generated consisting of various hydroxylated molybdenum oxide fluorides. The list of possible intermediates makes it difficult to assign specific bonds to the bands observed between  $500\text{--}1000\text{ cm}^{-1}$  during the intermediate steps of the hydrolysis reaction. In Particular, the peak at  $850\text{ cm}^{-1}$  is difficult to assign and does not align with reported Mo-F or Mo=O stretching. Bridged Mo-O stretching has been attributed to this region for molybdenum oxides<sup>28</sup>, but the band appears to grow in before the formation of molybdenum oxides in our experiment. Future computational studies may contribute significantly to the elucidation of possible intermediate species during the reaction.

Previous literature<sup>65, 66</sup> has reported Mo=O stretching frequencies for  $\text{MoOF}_4$  to be between  $1030\text{--}1048\text{ cm}^{-1}$  and  $\text{MoO}_2\text{F}_2$  Mo=O frequencies to be around  $1009\text{ cm}^{-1}$ . These numbers align well with the  $1000\text{ cm}^{-1}$  band observed in the reaction, but the Mo-F stretching frequencies of these compounds are typically observed in the  $700\text{--}750\text{ cm}^{-1}$  range. The Mo-F stretching bands in this experiments appear at  $650\text{ cm}^{-1}$ , which is closer to the frequencies observed in  $\text{MoF}_5\text{O}^-$  and fluorine bridged  $\text{Mo}_2\text{F}_9\text{O}_2^-$  compounds. The feature observed

around  $750\text{ cm}^{-1}$  is in line with previously reported hydroxylated molybdenum oxides and also with fluorine bridged compounds<sup>25, 64-68</sup>.

The spectra initially show the reactant species modes, then  $\text{MoF}_6$  and  $\text{H}_2\text{O}$  associated modes begin to disappear, and bands attributed to  $[\text{MoO}/\text{MoO}][\text{H}_{\text{net}}]$  are generated. Next, the bands attributed to  $[\text{MoO}/\text{MoO}][\text{H}_{\text{net}}]$  disappear as bands associated with  $\text{MoO}_3$  appear. Finally, a band attributed to the formation of  $\text{MoO}_3$  hydrate particulates grows in. The progression of the bands within the  $500\text{--}1000\text{ cm}^{-1}$  region and the depletion and subsequent regeneration of defined OH stretching bands leads to the proposed reaction pathway presented in Scheme 1.

Scheme 1. Proposed reaction for condensed phase  $\text{MoF}_6$  hydrolysis.

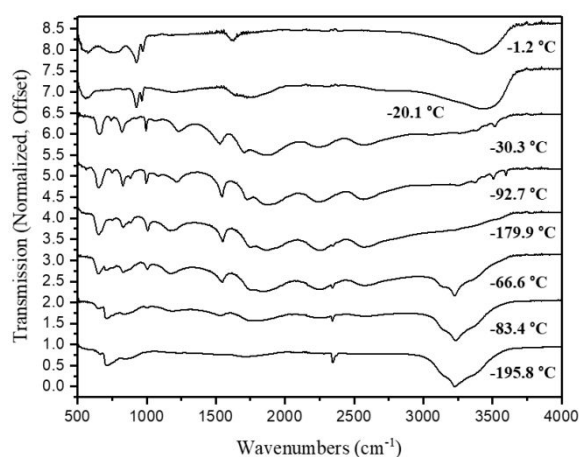
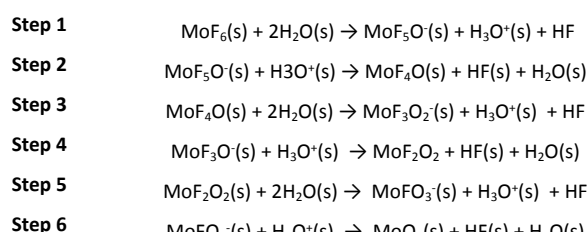


Figure 4 FTIR spectrum of the  $\text{MoF}_6$  hydrolysis while refreezing the sample. Liquid nitrogen was added to the cryocell to refreeze the sample in between each spectrum. During this process, the sample was allowed to cool down to  $-195.8\text{ °C}$  and warm back up to the labelled temperatures where the reaction proceeded.

The proposed reaction steps explain the appearance of the broad continuous aqueous proton bands which appear and is in line with previously reported hydrolysis products of  $\text{MoF}_6$ . Since the experiment was performed under vacuum, HF and excess water were likely pulled off and are not present in the final product.

#### Addition of $\text{CO}_2$ Insulating Layer

The experiment was also conducted using a  $\text{CO}_2$  insulating layer between  $\text{MoF}_6$  and  $\text{H}_2\text{O}$ , Figure 3. The insulating layer was added to ensure the reaction between the  $\text{MoF}_6$  and water could not proceed at liquid nitrogen temperatures. Although this experiment confirms that an insulating layer is not necessary to prevent the hydrolysis reaction under the non-insulated experimental conditions, the subsequent reaction pathway has several noticeable differences from the initial experiment. The differences in the two experiments could be attributed to the increased temperature of the reactants upon initial contact. In the non-insulated experiment, the Boltzmann distribution of the thermal energy of the reactants results in many of the steps of the reaction overlapping. In the experiment with the insulating layer, all the reactants have been brought up to the temperature required to boil off  $\text{CO}_2$  (approx.  $-140^\circ\text{C}$  under vacuum conditions) before interaction begins. As a result, most of the reactants in the experiment with the insulating layer have enough energy for the reaction to proceed as soon as the insulating layer has been removed. The reaction proceeds in a much clearer, stepwise fashion which considerably reduces the number of overlapping bands in the  $500 - 1000\text{ cm}^{-1}$  region.

The HF stretching mode around  $3600\text{ cm}^{-1}$  behaves much more consistently with the proposed reaction pathway. The  $3600\text{ cm}^{-1}$  band is initially absent and grows in, starting weak and broad before increasing in intensity and becoming sharper over time. The HF is likely being pulled off by the vacuum

conditions of the experiment before the defined OH stretching water bands begin to grow in. Since HF is a by-product of the initial reaction step the appearance of an HF band is expected quite early in the reaction. HF ingrowth is predicted to run concurrently with the initial depletion of the water bands. The hydrolyzing of the fluorine is observed by the development of the aqueous proton network, which is the result of HF being dissolved in the amorphous solid water network<sup>37</sup>. As the reaction progresses, the HF concentration reaches a point where the HF stretching mode can be detected. The excess water and HF is eventually boiled off and water is regenerated in the final step of the reaction forming a  $\text{MoO}_3$  hydrate complex.

The band around  $850\text{ cm}^{-1}$  that appears concurrently with the assigned  $[\text{Mo}=\text{O}/\text{MoO}][\text{H}_{\text{net}}]$  stretching mode in the non-insulated experiment is significantly suppressed in the insulating layer experiment, as well as many of the minor peaks within the  $500-1000\text{ cm}^{-1}$  range. Assignment of the bands during the intermediate portion of the reaction between  $500-1000\text{ cm}^{-1}$  was difficult in the original experiment due to the large number of possible intermediate species. These bands were attributed primarily to the formation of bridged compounds. The absence of these bands in the insulating layer experiment suggests they are not the result of simple sequential hydrolysis, but rather due to some intermediate or dimer complex formation which is either not forming or has a lifetime too fast to resolve under the reaction conditions created by adding an insulating layer of  $\text{CO}_2$ .

#### Refreezing Experiment

Refreezing of the reaction was done by addition of liquid nitrogen to the Dewar and re-cooling the cold finger after the reaction had begun, Figure 4. Re-cooling the cold finger "froze" the reactions in place, preventing the reaction from proceeding and sharpening many of the bands associated with the intermediate steps. The refreezing of the reaction had no impact on the formation of the final product. The intermediate

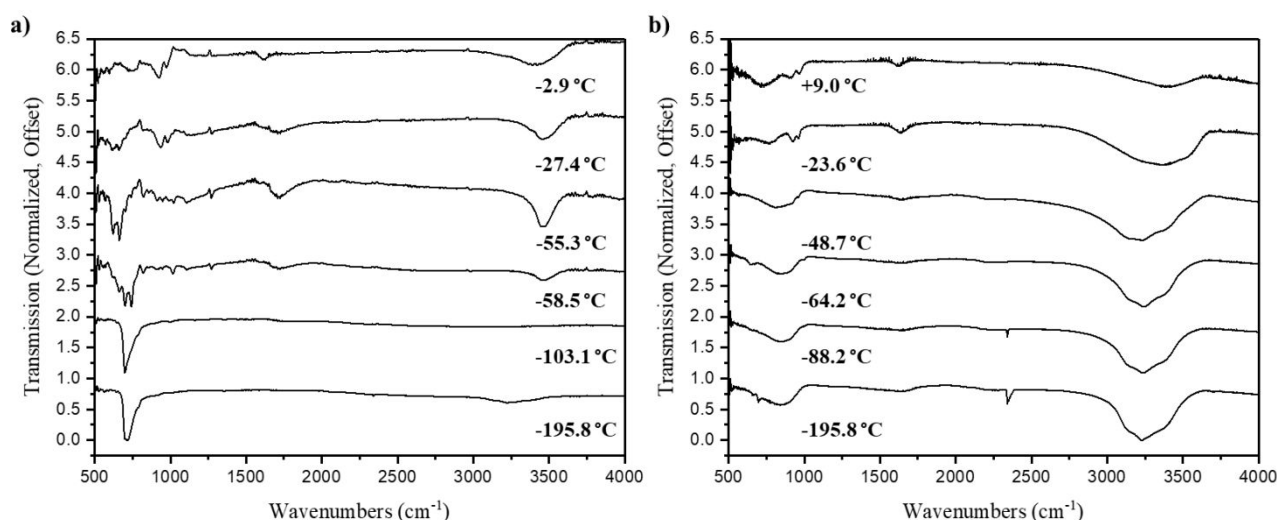


Figure 5 FTIR spectrum of the  $\text{MoF}_6$  hydrolysis under different limiting reactant conditions. a) limiting the amount of water layered on top of a thick layer of  $\text{MoF}_6$  layer, and b) limiting the amount of  $\text{MoF}_6$  initially layered followed by a thick layer of air to ensure excess water is present for the reaction. For each condition, reactants were allowed to slowly warm to room temperature while the IR spectrum was monitored.

bands between  $500\text{ cm}^{-1}$  and  $1000\text{ cm}^{-1}$  can be captured with the minor peak and shoulder between  $650\text{ cm}^{-1}$  and  $850\text{ cm}^{-1}$  appearing as they did in the original experiment. Both the minor peak and shoulder grow in around the same time, but the shoulder peak around  $850\text{ cm}^{-1}$  is completely depleted before the minor peak around  $650\text{ cm}^{-1}$  disappears. In addition, multiple sharp peaks appear in the HF stretching region, suggesting the HF produced is not trapped in a uniform network.

### Effect of Reactant Concentrations

The relative ratio of  $\text{MoF}_6$  and water were varied to assess the impact of limiting reactants on the overall reaction path and product production, Figure 5. Limiting the amount of water did not affect the formation of the final product and the general progression of the reaction is also unaltered. Therefore, the conclusion was drawn that the extended hydrogen bonding and aqueous proton networks are not required for the reaction to proceed to completion, which should be true since gas phase hydrolysis of  $\text{MoF}_6$  is possible. Interestingly, the final product is spectrally different when  $\text{MoF}_6$  is the limiting reactant. In the  $\text{MoF}_6$ -limited case, the  $750\text{ cm}^{-1}$  band attributed to the aggregation of  $\text{MoO}_3$  hydrates appears earlier and is much broader in the final product while the  $650\text{ cm}^{-1}$  band does not appear or is obscured. The change in these two bands must be attributed to the large excess of water still around during the formation of the  $\text{MoO}_3$ . In the previous experiments, all the well-defined OH stretching has disappeared before the transition from molecular  $\text{MoO}_3$  hydrate to  $\text{MoO}_3$  hydrate particulates. This suggests that excess water was not being pulled off by vacuum, but instead water was acting as the limiting reactant of the hydrolysis and was being completely consumed. When  $\text{MoF}_6$  is the limiting reactant, the excess water appears to have facilitated particulate aggregation.

### Conclusions

The work described herein reports a detailed spectroscopic study of the hydrolysis reaction of  $\text{MoF}_6$ . The reaction intermediates were demonstrated to be trapped in the disordered H-bonded network of a cryogenic amorphous thin film formed by sequential deposition of  $\text{MoF}_6$  and  $\text{H}_2\text{O}$  followed by heating. The hydrolysis reaction does not require the presence of the H-bonded network to proceed, but the relative abundance of  $\text{H}_2\text{O}$  does affect the condensed phase particulate formation. An insulating layer can be used to separate the two reactants but is not necessary to prevent the spontaneous hydrolysis of  $\text{MoF}_6$  at liquid nitrogen temperatures. A simple reaction pathway was proposed for the hydrolysis of  $\text{MoF}_6$  based on the available spectroscopic data and product identification.

Even though the final product of this reaction is not identical to the corresponding  $\text{UF}_6$  hydrolysis reaction, understanding the hydrolysis process for  $\text{MoF}_6$  can lead to more informed predictions for  $\text{UF}_6$  hydrolysis. The work also demonstrated the ability for the condensed phase hydrolysis of  $\text{MoF}_6$  to result in

particulate formation and for the particulates to be a function of the available water. Assuming similar pathways of particulate formation for  $\text{UF}_6$ , it would be expected that condensed phase particulates can also form.

### Experimental

#### Cryogenic Layering

A cryogenic cell was custom designed at Savannah River National Laboratory to perform experiments at liquid nitrogen temperatures. The cell was held at vacuum and was made with 2-inch diameter uncoated zinc selenide windows from Edmund Optics in order to view the samples, which were deposited onto a 18 mm diameter,  $150\text{ }\mu\text{m}$  thick polished polycrystalline diamond window from Applied Diamond, Inc.. A diamond surface was chosen as a substrate due to its transparency in the infrared and excellent thermal conductivity. The substrate was mounted on a copper cold finger extended from an open liquid nitrogen Dewar. The temperature of the cold finger was controlled by liquid nitrogen in the Dewar as the cold finger could be kept at liquid nitrogen temperatures by the addition of liquid nitrogen to the Dewar or warmed by allowing the liquid nitrogen to evaporate and exposing the copper cold finger to room temperature. The cryogenic cell was attached to a volume calibrated gas manifold which was used to control the amount of gas condensed onto the surface of the substrate. Thin films could be deposited onto the clean diamond surface after it has been cooled to  $-198\text{ }^\circ\text{C}$  by backfilling the chamber with gaseous samples admitted through a high-precision leak valve. Gaseous samples of  $\text{MoF}_6$  were admitted from high purity cylinders, purchased from Alfa Aesar, through the gas manifold, while atmospheric water and carbon dioxide were admitted via a controlled air leak onto the substrate. The layers were generated separately to calibrate the gas flow from the manifold to the cryogenic cell and control the thickness of the individual layers (S2). Once the samples were layered on the diamond surface, reactions were monitored in a Jasco FT-IR 6300 (Jasco Inc, Easton, MD, USA) while the layers were allowed to heat to room temperature over the course of 4 to 5 hours. A scan was taken every 30 seconds with a  $2\text{ cm}^{-1}$  resolution and a 400 to  $6000\text{ cm}^{-1}$  spectral range. The cold finger temperature was monitored using a thermocouple mounted to the end of the copper rod and digitally recorded over the course of the reaction using a DAC and LabVIEW program.

#### UV/Vis

The product was collected from the diamond substrate via a cotton swap and dissolved in D.I. water. The resulting liquid was then analysed with a custom-built UV – Vis using a xenon lamp light source.

#### Scanning Electron Microscopy

In order to collect product to be analysed with scanning electron microscopy (SEM), small carbon wafers were placed at the bottom of the cryogenic cell and a standard reaction was set up and executed. The wafers were then collected and analysed with a JEOL 2010 TEM and Hitachi SU8200 SEM.



## Conflicts of interest

There are no conflicts to declare.

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