HSO$_3$Cl: A Prototype Molecule for Studying OH-stretching Overtone Induced Photodissociation

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Abstract

Vibrationally induced photodissociation in sulfurochloridic acid (HSO$_3$Cl) is found to be a viable process to form SO$_3$ and HCl from excitations of the OH-stretching overtone starting at $\nu_{\text{OH}} = 4$. Reactive molecular dynamics simulations on a fully-dimensional potential energy surface fitted to MP2 calculations show that hydrogen transfer and HCl elimination compete with one another on the nanosecond time scale. For excitation with 5 and 6 quanta in the OH-stretch direct elimination of HCl is the dominant process on the several hundred picosecond time scale. At longer times, HCl formation is preceded by intramolecular hydrogen transfer and concomitant excitation of torsional degrees of freedom. As HSO$_3$Cl is a suitable proxy for H$_2$SO$_4$, which is relevant for weather and climate in the upper atmosphere, it is concluded that vibrationally induced photodissociation is a possible mechanism for H$_2$SO$_4$ decomposition. Final state energy distributions for different internal degrees of freedom are predicted which should be observable in laboratory measurements.
I. INTRODUCTION

The photochemistry of halogen- and sulfur-containing compounds which are involved in acid rain and aerosol formation in the tropo- and stratosphere, are of significance to the chemistry of the atmosphere. This led to extensive studies, both experimental and theoretical, concerning a wide range of chemical reactions relevant for climate and weather.\(^1\) In 1982 Hofmann and Rosen suggested that the formation of condensation nuclei from sulfur compounds is dominated by photochemical processes and they related the periodic de- and increase of these particles to the cycle of polar night and day around the poles.\(^2\) This seasonal change in the solar radiation affects directly the stratospheric aerosol layer by altering the rate of photochemical reactions of sulfur-containing compounds and thereby influences the climate and weather.

One of the central sulfur compounds in the stratospheric aerosol layer is sulfuric acid (H\(_2\)SO\(_4\)).\(^2\)\(^-\)\(^4\) Sources for H\(_2\)SO\(_4\) in the atmosphere are generally sulfur-containing compounds which are primarily emitted as dimethyl sulfide (DMS) and sulfur dioxide (SO\(_2\)).\(^5\)\(^-\)\(^9\) These compounds are oxidized in the atmosphere to form sulfur trioxide (SO\(_3\)) and subsequently hydrolyzed to H\(_2\)SO\(_4\).\(^10\)\(^,\)\(^11\) Among other, stratospheric sulfate aerosol has a global dimming effect in the atmosphere and increases the number of aerosol particles which can limit the effect and impact of global warming.\(^12\) This makes H\(_2\)SO\(_4\) a suitable candidate in climate engineering projects.\(^13\) Additionally, the formation of a layer of dispersed aerosol found at high altitudes (between 15 to 35 km) which influences stratospheric ozone depletion, the so called Junge layer\(^14\), can be explained by the cycling between SO\(_2\) and H\(_2\)SO\(_4\).\(^15\)\(^-\)\(^17\)

Sulfur-containing compounds, in particular the interplay between H\(_2\)SO\(_4\) and SO\(_2\), are important in the formation of atmospheric aerosols. The observed anomalous enhancement in particle concentration during springtime and the finding that SO\(_2\) concentration from high altitude photolysis of H\(_2\)SO\(_4\) is constant or increases with altitude\(^16\) suggested that at high altitudes photolysis of sulfuric acid is the source of SO\(_2\).\(^18\) However, experimental and computational investigations were unable to find the electronic absorption spectrum of H\(_2\)SO\(_4\) down to 140 nm.\(^19\)\(^-\)\(^23\) According to Lane and Kjaergaard\(^24\) the lowest calculated electronic transition is at 139 nm. Hence, UV excitation from solar photons is improbable to produce SO\(_2\) from H\(_2\)SO\(_4\) and an alternative reaction mechanism for H\(_2\)SO\(_4\) decomposition was required.
Experimentally and computationally it was previously found that vibrational excitation can induce bond-selective photodissociation.\textsuperscript{25–27} It was therefore proposed that highly-excited OH-stretching states can lead to dissociation of $\text{H}_2\text{SO}_4$.\textsuperscript{18} The vibrational spectrum of $\text{H}_2\text{SO}_4$ in the near-infrared and visible regions has been intensely studied, both experimentally\textsuperscript{19,28–31} and computationally.\textsuperscript{22,23,31–33} Nevertheless it was not possible to experimentally verify that vibrational overtone-induced photodissociation is operative in $\text{H}_2\text{SO}_4$. This is due to particular experimental difficulties involving this species. By mixing known vapor flows of $\text{H}_2\text{O}$ and $\text{SO}_3$ at elevated temperature (400-500 K) $\text{H}_2\text{SO}_4$ can be generated in the vapor phase ($\text{H}_2\text{O} + \text{SO}_3 \rightarrow \text{H}_2\text{SO}_4$).\textsuperscript{19} However, $\text{H}_2\text{SO}_4$ has an extremely low vapor pressure, and exists in equilibrium with $\text{H}_2\text{O}$ and $\text{SO}_3$ in the gas phase. Therefore this approach leads to difficulties in distinguishing the reactants and products.\textsuperscript{29,34}

Because of the above mentioned difficulties and the widespread relevance of the atmospheric decomposition of sulfates, sulfuric acid derivatives have been considered to provide experimental evidence for characterizing OH-stretching induced dissociation in the gas phase.\textsuperscript{29,34,35} Such candidate compounds are sulfurochloridic acid (HSO$_3$Cl) and sulfurofluoridic acid (HSO$_3$F) which are related to $\text{H}_2\text{SO}_4$\textsuperscript{29,34}, but experimentally more amenable. The dissociation energy to eliminate HCl and HF from HSO$_3$Cl and HSO$_3$F, respectively, are similar to those of $\text{H}_2\text{O}$ from $\text{H}_2\text{SO}_4$ (32-40 kcal/mol).\textsuperscript{34} Using a simple one-dimensional anharmonic oscillator local model, the overtone spectra of sulfuric acid derivatives have been computed, which found that excitation of the OH-stretching transition with $\Delta v \geq 4$ and $\Delta v \geq 5$, will provide sufficient energy for photodissociation of HSO$_3$Cl and HSO$_3$F, respectively.\textsuperscript{29,34,35} Finally, \textit{ab initio} molecular dynamics (MD) simulations have demonstrated that the decomposition of these derivatives, similar to what has been proposed to $\text{H}_2\text{SO}_4$, is possible \textit{via} excitation of higher OH-stretching overtone transitions.\textsuperscript{29,34} At the semi-empirical PM3 level, it was found that for HSO$_3$Cl, excitation of $v \geq 4$, or $v \geq 5$ (PM3) can induce the elimination reaction.\textsuperscript{34} For photodissociation of HSO$_3$F excitation of the $v \geq 7$ OH-stretching state is necessary with PM3, whereas excitation of the $v \geq 5$ overtone is sufficient at the MP2/TZP level.\textsuperscript{34} The average reaction time for both, HSO$_3$Cl and HSO$_3$F, was of the order of 10 ps, determined from 32 trajectories.\textsuperscript{34} However, previous reactive MD simulations on $\text{H}_2\text{SO}_4$ have demonstrated that for converged reaction time distributions several thousand trajectories are required.\textsuperscript{32,33}
In order to characterize and understand the dynamics of vibrationally induced decomposition of HSO$_3$Cl and to guide future experimental efforts, Multi-Surface Adiabatic Reactive Molecular Dynamics (MS-ARMD) simulations were carried out. This method is capable of treating multiple reaction pathways based on force fields to describe the adiabatic potential energy surface and the classical reaction dynamics of the system, and to run and analyze a statistically significant number of trajectories. Here, the dynamics on a fully dimensional reactive potential energy surface (PES) capable of following intramolecular H-transfer and HCl elimination in HSO$_3$Cl (see Figure 1) is investigated. This provides the information to analyze the yields and determine rate coefficients along with the final state energy distribution in the products as a function of the vibrational overtone that was initially excited. This information can be directly compared with forthcoming experiments.

FIG. 1: Graphical representation of the two possible reaction pathways involving HSO$_3$Cl considered here. Top trace: intramolecular H-transfer (O$_A$H $\rightarrow$ O$_B$H), bottom trace: HCl elimination.
II. REACTIVE MOLECULAR DYNAMICS SIMULATIONS

The global PES for the reactions of HSO$_3$Cl is based on reference electronic structure calculations at the MP2/6-311G++(2d,2p) level of theory. All ab initio calculations were performed with Gaussian09\textsuperscript{37}. The energy at the equilibrium geometry of HSO$_3$Cl was chosen as the global zero of energy. The functional form of the force field which describes the HSO$_3$Cl, SO$_3$ and HCl states, including the SO$_3$···HCl van der Waals complex has the following form:

\[
V(x) = \sum_{\text{bonds}} D_e (1 - e^{-\beta(r-r_0)})^2 + \sum_{\text{angles}} K_\theta (\theta - \theta_0)^2
+ \sum_{\text{Urey-Bradley}} K_{UB} (S - S_0)^2 \sum_{\text{dihedral}} H_\phi (1 + \cos(n\phi - \delta))
\]

\[
+ \sum_{\text{impropers}} K_\omega (\omega - \omega_0)^2
+ \sum_{ij} \left\{ \frac{n \varepsilon_{ij}}{m - n} \left[ \left( \frac{r_{\min,ij}}{r_{ij}} \right)^m - \frac{m}{n} \left( \frac{r_{\min,ij}}{r_{ij}} \right)^n \right] + \frac{q_i q_j}{4\pi\varepsilon_0 r_{ij}} \right\}
\]

where $r_0$ is the equilibrium bond distance, $D_e$ is the dissociation energy, and $\beta$ controls the steepness and width of the Morse potential.\textsuperscript{36} Additionally a generalized Lennard-Jones potential ($\sum_{ij}$) is included, in order to better represent the van der Waals interaction between atoms in the reactive region.\textsuperscript{36} The PESs of the three molecules (HSO$_3$Cl, SO$_3$ and HCl) together with the van der Waals complex (SO$_3$···HCl) were parameterized independently. Specifically, the HSO$_3$Cl state is described by all terms in equation 1, whereas for SO$_3$ only bonds, angles, Urey-Bradley, improper dihedrals and non-bonded terms, are needed. For HCl a bonded term and non-bonded terms (for the van der Waals complex) were required. For the reactive PES of the overall system the individual PESs were joined into a global reactive PES using the MS-ARMD switching method and were combined with Gaussian $\times$ polynomial (GAPOs) functions which provide the necessary flexibility to model the barrier crossing region.\textsuperscript{36} The interpolation of the PESs is accomplished through energy-dependent weights $w_i(E)$.\textsuperscript{36}

Details of the fitting are described in the Supplementary Information. Optimization of the force field yields parameters for the global reactive PES which allows to follow the H-transfer and HCl elimination reactions. The root-mean-square deviations (RMSD) between ab initio reference energies and FF energies are 0.22 and 0.29 kcal/mol, for the barrier region of HCl elimination and intramolecular H-transfer, respectively, see red curves in Figure 2 A and B.
FIG. 2: Minimum Energy Path for A) HCl elimination and B) intramolecular H-transfer. Black circles represent the *ab initio* reference energies, red curves represent the FF energies including GAPOs.

The RMSD between the energy-minimized structures at the MP2/6-311G++(2d,2p) level and using the force field is 0.004 Å whereas for the SO$_3$····HCl van der Waals complex the RMSD is 0.053 Å. A comparison of equilibrium bond lengths, angles and dihedral angles is provided in Table S-I (Supplementary Information). The geometries in Table S-I are also in good agreement with those computed using MP2/TZP (Dunning’s triple-ζ+ polarization). In addition, a normal mode analysis was performed in order to compare the experimentally observed vibrational frequencies with those from the MP2 and FF calculations, see Table S-II (Supplementary Information). The average differences between experimental, MP2 and FF harmonic frequencies for HSO$_3$Cl are 4.56 and 4.71 cm$^{-1}$, respectively which establishes the quality of the force field.

Particularly relevant for the reaction rates are the barrier heights. The MP2/6-311G++(2d,2p)
computed values are close to those determined previously using CCSD(T)/aug-cc-pV(T+d)Z. For HCl elimination the value of 34.0 kcal/mol compares with 33.1 kcal/mol from the present study. As the final global PES can describe in good agreement the barrier height from ab initio methods, and the geometry in the equilibrium state is well represented for the FF, the reactive PES is considered to be parameterized in a meaningful way. Contrary to an ab initio MP2/6-311G++(2d,2p) single point calculation, the reactive FF requires a fraction of the computational cost, which is highly relevant for dynamical studies.

The quality of the reactive PES is shown in Figure 3. It reports the energies of the reference MP2 calculations compared with the FF. For the representative structure in thermal equilibrium (300 K), the IRCs (HCl elimination and intramolecular H-transfer, obtained from individual scans) and HCl + SO₃ complex, the absolute average difference between all the reference MP2 energies and all the FF is 0.52 kcal/mol.

All MD simulations were carried out with CHARMM with provisions for bond-breaking and bond-formation through MS-ARMD. Starting from a geometry optimized structure of HSO₃Cl, the system was heated to 300 K. The equations of motion were propagated with the leapfrog Verlet algorithm with a time step of \( \Delta t = 0.1 \) fs during 40 ps and equilibrated for 40 ps, followed by 250 ns of free dynamics simulations (i.e. constant energy NVE), where a structure was collected every 50 ps. The small time step is required to appropriately follow the rapid dynamics of the H-atom.

The nonequilibrium state was prepared by scaling the instantaneous velocity vector at the moment of excitation along the normal mode of interest which is the \( \nu_{OH} \) mode in the present case. Similar to the situation in \( \text{H}_2\text{SO}_4 \) this mode has predominantly local OH-stretching character. Such a scheme, which modifies only the kinetic energy, has already been successfully employed for \( \text{H}_2\text{SO}_4 \) and for studying proton transfer in small H-bonded complexes. An alternative approach is to change both, kinetic and potential energy, by modifying positions and velocities. While changing only the kinetic energy can lead to accelerated redistribution of vibrational energy, modification of positions and momenta requires a short equilibration period to avoid artifacts due to close proximity of atoms which may be a potential drawback in this approach. Nonequilibrium excitation of a thermal ensemble as described above is closest to the situation...
encountered in a laboratory experiment and was therefore preferred over directly generating quasiclassical (QCT) nonequilibrium initial conditions.

The OH stretch potential can be realistically described by a Morse potential for which the exact energy levels are known. The excitation energies for $\nu_{\text{OH}} = 3, 4, 5, \text{and } 6$ are: $11699 \text{ cm}^{-1}$ ($33.3 \text{ kcal/mol}$), $14760 \text{ cm}^{-1}$ ($42.2 \text{ kcal/mol}$), $17662 \text{ cm}^{-1}$ ($50.5 \text{ kcal/mol}$), and $20405 \text{ cm}^{-1}$ ($58.3 \text{ kcal/mol}$) using $D_e = 122.88 \text{ kcal/mol}$ and $\beta = 2.11 \text{ Å}^{-1}$ from the current O-H bond in the MS-ARMD potential. An alternative means to select initial conditions is described in the Results section.

III. RESULTS

From 5000 independent trajectories with a maximum simulation time of 2.5 ns, more than 90% of the trajectories were reactive when $\nu_{\text{OH}} = 5$ or 6 are excited (see Figure 4). Both processes, (I) HCl elimination without prior H-transfer (dashed line), called direct, and (II) HCl elimination preceded by intramolecular H-transfer (dotted dashed line), called indirect, are observed. Direct HCl elimination is found for 0.2, 10.4, 28.5, and 42.8% of the cases for $\nu_{\text{OH}} = 3, 4, 5, \text{and } 6$. HCl elimination preceded by intramolecular H-transfer was observed for 4.2, 62.0, and 54.9% of the cases for $\nu_{\text{OH}} = 4, 5$ and 6 whereas for $\nu_{\text{OH}} = 3$ this reaction channel was not observed. Intramolecular H-transfer without HCl elimination within 2.5 ns (leading to complete intramolecular vibrational energy redistribution (IVR), which strongly decreases the decay probability) was found for 10.2, 35.3, 7.3, and 0.08% of the cases for $\nu_{\text{OH}} = 3, 4, 5, \text{and } 6$.

For excitation with $\nu_{\text{OH}} = 5$ and 6 decomposition occurs on the hundred picosecond time scale. Overall 50% of the trajectories show HCl elimination within 500 ps for excitation with $\nu_{\text{OH}} = 6$. For $\nu_{\text{OH}} = 5$ the proportion is around 34% after 500 ps. The majority of these trajectories that show decomposition for $\nu_{\text{OH}} = 5$ and 6 lead to direct HCl elimination, without previous H-transfer. On a longer time scale indirect elimination becomes dominant, as some time is required for the "hydrogen hopping". For $\nu_{\text{OH}} = 3$ and 4 the decomposition rates are still linear and far from stationary which would require much longer simulation times, particularly for the excitation with 3 quanta. It is worthwhile to mention that the mean collision time was estimated to be around 130
FIG. 3: Quality of the reactive PES. The reference data points are calculated at the MP2/6-311G(d,p) level of theory and compared to the parameterized surface.

However, multiple 1000 simulations on this time scale become computationally prohibitive even with an empirical force field. The effect of deuteration was assessed by running 5000 trajectories for DSO₃Cl with νOD = 5 which corresponds to 13546 cm⁻¹ (36.2 kcal/mol). Within 2.5 ns, 2% of these trajectories showed DCl elimination, nearly all of them following process I. Hence, for the deuterated species much higher vibrational excitations are required.
FIG. 4: Percentage of HCl elimination as a function of simulation time. Top panel: Total number of reactive events (direct HCl elimination + HCl elimination preceded by intramolecular H-transfer). Green, blue, red and black curves correspond to excitation of $\nu_{OH} = 3$, 4, 5 and 6, respectively. The red dashed curve corresponds to DCl elimination in DSO$_3$Cl with $\nu_{OD} = 5$. Bottom panels: Total number of reactive events (solid), direct HCl elimination (dashed), and HCl elimination preceded by intramolecular H-transfer (dotted dashed) for $\nu_{OH} = 3$, 4, 5 and 6 in the same color code as in the top panel.

for photodissociation which leads to a smaller cross section due to reduced overlap of the initial and final vibrational states than for the hydrogenated species.

Compared with a previous study on H$_2$SO$_4$ the considerable differences in the number of reactive events for excitation with 5 or 6 quanta are found. For HSO$_3$Cl the percentage of trajectories which lead to HCl elimination (yield) is $\approx 60\%$ for $\nu_{OH} = 5$ after 1 ns, which is markedly increased compared to 25\% in H$_2$SO$_4$ for the same OH stretching overtone within 1 ns, even though the barrier in H$_2$SO$_4$ is slightly lower. This can be explained by the fact that the excitation
energy in HSO\textsubscript{3}Cl is higher. For $\nu_{\text{OH}} = 5$ the difference between H\textsubscript{2}SO\textsubscript{4} and HSO\textsubscript{3}Cl is 3.3 kcal/mol. Furthermore, for sulfuric acid direct elimination was the dominant process for all excitation levels. Only 3.4% and 12.2% of the trajectories in H\textsubscript{2}SO\textsubscript{4} for 5 and 6 quanta, respectively, were found to proceed via indirect elimination.\textsuperscript{33} This can be explained by the fact that in H\textsubscript{2}SO\textsubscript{4} direct elimination occurs more rapidly than in HSO\textsubscript{3}Cl. The maximum lifetime for, e.g. $\nu_{\text{OH}} = 5$ is 300 ps in H\textsubscript{2}SO\textsubscript{4} compared to 500 ps in HSO\textsubscript{3}Cl. Furthermore the rate of intramolecular H-transfer is constant in time in H\textsubscript{2}SO\textsubscript{4}, which can be explained by the fact that in H\textsubscript{2}SO\textsubscript{4} a transformation of symmetry, and therefore a change in the energy, occurs. In HSO\textsubscript{3}Cl the intramolecular H-transfer is symmetric.

Most relevant for future experiments is the final state analysis of the products, which is an observable.\textsuperscript{43} In order to characterize the energy distribution after vibrational excitation and subsequent HCl elimination, the translational, rotational and vibrational energies in the reaction products were determined following established methodologies, as described in the Supplementary Information. The analysis was carried out for all trajectories that experienced HCl elimination within 2.5 ns for $\nu_{\text{OH}} = 4, 5,$ and 6. The analysis for $\nu_{\text{OH}} = 3$ was excluded due to the small number of reactive events. Final state analysis starts after HCl elimination when the sulfur-chlorine distance exceeds 20 Å. Trajectories were then continued for an additional 5 ps during which 500 snapshots for each trajectory were collected and analyzed. Translational, rotational, and vibrational energies as well as the total angular momentum were computed for every snapshot. Rotational and vibrational energies were averaged over the 500 snapshots. From these averages the probability distributions of the averaged values were determined.

Figure 5 shows the distribution of translational, rotational, and vibrational energies of the products for $\nu_{\text{OH}} = 4$ (blue), 5 (red), and 6 (black). The peak position of translational energies differ by less than 3 kcal/mol for the three excitation levels. For SO\textsubscript{3} a maximum peak for excitation with all quanta is found at about 5 kcal/mol. Rotational energy distributions for HCl peaks at about 3.5 kcal/mol (corresponding to a maximum angular momentum of $j = 11$), with a difference of the peak position of less than 1 kcal/mol. For SO\textsubscript{3} the rotational energy shifts towards higher energies for lower $\nu_{\text{OH}}$. The vibrational energies show similar behavior for HCl and SO\textsubscript{3}. In both cases the vibrational energy distribution widens for higher overtone excitation. For SO\textsubscript{3} the maximum peaks at around 11, 15, and 19 kcal/mol for $\nu_{\text{OH}} = 4, 5$ and 6 respectively.
The $j-$distribution (Figure S-III (Supplementary Information)) for SO$_3$ exhibits a single maximum around $j = 60$ to 70 for all investigated excitations. Contrary to that, for HCl a slight bimodal $j-$distribution for higher vibrational excitations, specifically for $v_{OH} = 6$ is found. The two peaks can be assigned to the direct and indirect product channels. Direct elimination, with a peak around $j = 10$, is associated with higher vibrational energy, whereas indirect elimination leads to a higher $j-$value (around $j = 12$).

An alternative method to characterize final states from MD simulations, in particular for vibrational degrees of freedom, is the Gaussian binning procedure$^{44}$ This was applied to the diatomic product as this would also be most relevant from an experimental perspective. Gaussian binning consists of assigning larger weights to final states with fractional vibrational quantum numbers $x_i$ closer to an integer value $n_i$. For the present case of a diatomic and following Ref.$^{44}$ the weight is $w = \sqrt{\beta} e^{-\beta(x-n)^2}$ where $x$ and $n$ are the fractional and integer quantum numbers of HCl in the product state, respectively. The parameter $\beta$ should be chosen sufficiently large in order to reduce weights of final states near half-integer values and was $\beta = 20$ in the present case. Sebastian check: For excitation of HSO$_3$Cl with $v = 5$ the final state population for the HCl stretching mode is 43% ($v_{HCl} = 0$), 30% ($v_{HCl} = 1$), 18% ($v_{HCl} = 2$), and 9% ($v_{HCl} \geq 3$). This only differs marginally from binning all final states equally ($w = 1$) and supports the final state analysis discussed above.

In order to validate the results, an alternative method to generate initial conditions was employed. Instead of starting from an equilibrium ensemble at 300 K, energies were drawn from a Maxwell-Boltzmann distribution using Metropolis Monte Carlo (MC) sampling and initial conditions generated according to an algorithm developed by Schranz et al.$^{45}$ For each drawn energy $E$, spatial coordinates are selected from the final state of a Markov chain generated by a Metropolis-type procedure.$^{46}$ Then, random momentum vectors are generated from a Gaussian distribution for each atom. Finally, the momentum vectors are rescaled such that the total kinetic energy $T = E - V$, where $V$ is the potential energy of the generated configuration. For convenience, all generated conditions were transformed to a center-of-momentum frame. The potential energy distribution for the two different ensembles are similar to one another. The distribution of the potential energy for the ensembles generated from the MC sampling shows a

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FIG. 5: Final state analysis for HCl and SO$_3$. Blue, red, and black curves correspond to excitation of $v_{OH} = 4$, 5 and 6, respectively. The abbreviations trans., rot. and vib. stand for the translational, rotational, and vibrational energy, respectively.

maximum peak slightly closer to $\frac{15}{2}k_BT$, which is the thermodynamic value.

Figure 6 shows the decomposition probability for 1000 trajectories generated with the MC method. The excitation of the OH bond was applied in the same way as for the previous ensemble. Calculations with $v_{OH} = 3$ were excluded due to the small number of reactions found for the original set of initial conditions (see Figure 4). Qualitatively, the reaction rates are comparable to those from simulations in which excitation of the equilibrium ensemble were used, see Figure 4. Also, the relative importance of direct HCl elimination and elimination preceded by H-transfer to neighboring oxygen atoms is comparable. Quantitatively, the decomposition probability for all excitation is reduced by 25% to 30% compared to simulations with initial conditions taken from NVE simulations. Furthermore, the reaction slows down, as can be seen by comparing, e.g., the point at which excitation with $v_{OH} = 6$ reaches 50% reactivity. For the results shown in Figure 6
with the alternative set of initial conditions, this occurs at around 1 ns compared to \( \approx 500 \) ps for the original choice of initial conditions, see Figure 4. The same is found for the crossing point between direct and indirect elimination with \( \nu_{OH} = 5 \) and 6 which is shifted towards a longer times.

![Graph showing reaction kinetics](image)

**FIG. 6:** Reaction kinetics from 1000 trajectories with initial conditions from the Monte Carlo sampling. Top panel) Total number of reactive events (direct HCl elimination + HCl elimination preceded by intramolecular H-transfer). Blue, red and black curves correspond to excitation of \( \nu_{OH} = 4, 5 \) and 6, respectively. Bottom panels) Total number of reactive events (solid), direct HCl elimination (dashed), and HCl elimination preceded by intramolecular H-transfer (dotted dashed) for \( \nu_{OH} = 4, 5 \) and 6 in the same color code as in the top panel.

The trajectories run with the Monte Carlo initial conditions were analyzed in the same fashion as the original 5000 trajectories, see Figure S-IV (Supplementary Information). The energy distributions after HCl elimination are slightly shifted towards higher energies when compared to Figure 5. However, the distributions remain qualitatively similar. As for the vibrationally induced
decomposition reaction, conclusions drawn from the final state analysis are robust with respect to the two very different procedures to generate initial conditions for the non-equilibrium simulations.

FIG. 7: Probability distribution $p(r_{HO_A}, \phi)$, with $\phi$ as the dihedral angle between H-Cl-S-O$_A$, shown as contour lines. From left to right excitation with $v_{OH} = 4, 5,$ and 6. From top to bottom indirect trajectories showing H-transfer to both oxygen atoms, to O$_B$ only, and to O$_C$ only. The molecular structures represent the geometry of the region marked with a red cross (for each panel individually). $p(r_{HO_A}, \phi)$ was normalized to unity for each panel and the lowest isocontour is labelled in each case.

The decomposition reactions preceded by intramolecular H-transfer differ from direct HCl elimination primarily by the time scale on which dissociation occurs, see Figures 4 and 6. In order
to characterize the differences between these two reaction pathways, geometrical coordinates were sought which can be used to further analyze indirect elimination. Visual inspection of the dynamics suggested that the H-Cl-S-O_A dihedral angle $\phi$ could be a meaningful choice. The dihedral angle $\phi$ can distinguish between the three oxygen atoms in a symmetric way (with O_A on the mirror axis). The distance O_A helps to further distinguish the reactive trajectories, especially with respect to the position of the hydrogen to its acceptor. The trajectories can be grouped into those in which during the dynamics the hydrogen atom binds to oxygen atoms O_A and O_B, O_A and O_C, or to all three oxygen atoms.

Probability distributions $p(r_{HO_A}, \phi)$ from analyzing 1000 trajectories, taken from the ensemble in which excitation of the equilibrium was used, for each of the excitation levels are reported in Figure 7. Each column displays, from top to bottom, H-transfer to (I) both oxygen atoms, (II) O_B only, (III) O_C only before HCl elimination occurs. The columns report results for increasing number of OH-quanta from left to right. Each panel was individually normalized. For excitation with $\nu_{OH} = 4$ (left column) only a small number of trajectories show indirect HCl elimination: for cases (I) to (III) there are 2, 39 and 10 events, respectively. For excitation with 5 quanta these numbers increase to 295, 525, and 104 trajectories whereas for excitation with $\nu_{OH} = 6$ cases (I) to (III) occur for 356, 358, and 86 runs. Hence, for excitations with 4 to 6 quanta 90%, 10% and 20% of the trajectories, respectively, lead to HCl elimination without transfer between neighboring oxygen atoms.

The distribution of the angle $\phi$ reveals two states for each possible H-O_X (X = A, B, C) bond, see red crosses with corresponding molecular models in Figure 7. These states correspond to two synperiplanar positions of the hydrogen with respect to O_X. The probability distributions for the two states at $\phi = \pm 25^\circ$ and short HO_A bond length consist of the positions of the hydrogen atom sampled a) in the equilibrium trajectories of HO_A SO_2Cl before excitation and b) during the migratory dynamics after vibrational excitation. This explains the high probabilities in these two positions. The distribution of the H-O_A bond length broadens for higher vibrational excitations as the vibrational energy along the OH-bond increases (see left sides in the panels in Figure 7). With $\nu_{OH} = 4$ the hydrogen remains longer in a specific orientation since the HO_A S rotation barrier is rarely overcome. This – and the small number of trajectories in this class – leads to the asymmetric distribution along $\phi$ in the top left corner. For excitation with $\nu_{OH} = 5$ the distribution
$p(r_{HO_A}, \phi)$ for trajectories sampling all oxygen atoms is most symmetrical. This indicates that the dihedral barrier can be overcome and that extensive sampling of all oxygen positions by the hydrogen atom is possible before elimination. Finally, with $v_{OH} = 6$ the distribution in the top right corner is again less symmetric as the residence time of the hydrogen on each oxygen before elimination is shortened due to the high excitation energy. The middle column of Figure 7 illustrates the increasing H-O$_A$ amplitude as the number of quanta is increased from left to right. From these probability distributions it is evident that coupling between the OH-stretching vibration and the torsional degrees of freedom is an essential pathway for energy exchange which influences the rates and yields of HCl elimination on the picosecond time scale.

IV. CONCLUSION

In summary, the present work establishes that vibrational excitation of OH-overtones with 3 to 6 quanta leads to HCl elimination on the sub-nanosecond to ns time scale for HSO$_3$Cl. For short times (a few hundred ps) after vibrational excitation this process is usually direct, i.e. without previous intramolecular H-transfer and leads to characteristic energy distributions in the diatomic fragment which should be experimentally detectable. Direct and indirect HCl elimination occur whereby the latter is driven by appreciable excitation of torsional degrees of freedom and dominates at longer times. It is shown that the conclusions from the simulations are robust towards the particular choice of initial conditions. Experimentally corroborating such a reaction mechanism will be relevant for atmospheric chemistry at large, since under the conditions in the stratopause around the poles vibrationally excited states can survive sufficiently long for decomposition to occur. The reaction is fast (sub-ns to ns time scale) compared to collisionally induced quenching, which occurs on the hundred ns time scale.\textsuperscript{18,33} Hence, there is only little competition for the dissociative channel. The present work provides testable hypotheses for experiments and establishes a vibrationally driven photodissociation mechanism for HSO$_3$Cl which is a meaningful proxy for H$_2$SO$_4$ relevant in atmospheric chemistry.
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