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Vaibhav Singh, Chuan Cheng, Thomas Weinacht, and Spiridoula Matsika

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Quantum Contributions to Coulomb Explosion Imaging Revealed by Trajectory Surface Hopping Molecular Dynamics

Vaibhav Singh,[1] Chuan Cheng,[2] Thomas Weinacht,[2] and Spiridoula Matsika[1]

[1] *Temple University Department of Chemistry, Temple University, Philadelphia, PA 19121, USA*
[2] *Department of Physics, Stony Brook University, Stony Brook, NY 11790, USA*

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We compare trajectory surface hopping (TSH) calculations of the dissociation of quadruply charged formaldehyde molecules with basic classical calculations assuming Coulomb repulsion between singly charged atoms. The TSH calculations show very different dissociation dynamics from the classical Coulomb calculations, illustrating that Coulomb repulsion between point charges does not accurately describe the dissociation dynamics. By comparing the calculations with covariance measurements, we see much better agreement between the experiments and the TSH calculations, and evidence for the production of multiple electronic states of the tetracation via strong field ionization.

I. INTRODUCTION

Coulomb explosion imaging (CEI) has become a prominent technique to capture the instantaneous geometry of a molecule undergoing rapid structural changes. Hence, it has helped study the ultrafast dynamics of several molecules [1–6]. In this approach, multiple ionization in the electric field of an intense ultrafast laser pulse can build up positive charge in a molecule that causes the charged fragment ions to fly away due to Coulomb repulsion. Measuring the momenta of the fragments can then be used to recover the initial molecular geometry. This technique has helped provide structural and dynamical insight into an array of systems and processes, including the characterization of isomers[5–8], hydrogen migration[9–12], roaming[13] and the visualization of conical intersections[14]. The most common approximation used to model the phenomenon is to assume that every atom or fragment acquires a charge of +1 and Coulombic forces are the only forces acting between the fragments or atoms, resulting in Coulomb explosion [2, 3, 9, 15, 16]. Besides this simplistic approach, there are a few studies where *ab initio* dynamics, which include quantum effects such as bonding, non-adiabatic couplings, etc., have been used[12, 13, 17, 18]. However, the theoretical modeling in these studies either used a reduced dimensionality model, or only included the ground state of the charged species, or examined only partial fragmentation. The work of Endo *et al.*, which is closer to the present study, compared calculations for the dissociation of D_2CO^{3+} using classical Coulombic forces and the *ab initio* potential energy surfaces of D_2CO^{3+} in the ground state. While their work found reasonable agreement between the two approaches, two of the three particles involved are point charges, and it is not clear whether such agreement generally extends to more complicated fragmentation involving multi-electron fragment ions.

In this work, we try to understand the 4-body breakup of the deuterated formaldehyde tetracation (CD_2O^{4+}) due to Coulomb explosion using multireference electronic structure methods and on the fly nonadiabatic dynamics through semi-classical trajectory surface hopping (TSH)[19, 20] on multiple surfaces. To check if an elementary approach of a charge per atom could be simulated with the Coulombic Forces (CF), we compare the CF results with the non-adiabatic dynamics. We find significant differences between the two calculations, including the formation of metastable CO^{2+} in the TSH dynamics. Thus, we show that the approximation of one charge per fragment/atom and pure classical Coulomb forces is insufficient to accurately treat dissociation. Experimental work by Cheng *et al.*[21] showed some evidence of metastable CO^{2+}, which can be explained by the results described here. This work is also unique in terms of the direct comparison of CF and TSH results with the experimental results that sets another example of why quantum principles can be important in describing processes involved during CEI.

II. METHODOLOGY

A. Electronic Structure Methods

The geometry of neutral formaldehyde was optimized at the Density Functional Theory(DFT) [22, 23] level of theory using B3LYP[24–27] functional and 6-311+G(d)[28, 29] basis set. The potential energy surfaces (PESs) of CH_2O^{4+} were scanned along C-O stretch, and symmetric C-H stretch at 20 states averaged - complete active space self-consistent field (20SA-CASSCF(8,8))[30]/cc-pVDZ[31] level of theory using C_{2v} symmetry. Here, 5 singlets and 5 triplets were calculated separately for each of the four irreducible representations of C_{2v} symmetry, i.e., A_1, B_1, B_2, and A_2 summing up to 20 singlets and 20 triplets. A large number of states were included in order to identify the lowest energy states of the correct symmetry. Once that was done, the following discussion and dynamics focused on the first 6 singlet and 4 triplet states. The PESs of CO^{2+} singlets and triplets were also calculated at the 20SA-CASSCF(8,8)/cc-pvDZ

level of theory, averaging over five singlet and five triplet states for each irreducible representation in C_{2v} symmetry (which again is 20 singlets and 20 triplets in total). Here, the C_{2v} is used as a lower symmetry group to calculate the states of CO^{2+}, which has an actual symmetry of $C_{\infty v}$. The 3D PESs of CH_2O^{4+} were also calculated as a function of C-O stretch and symmetric C-H stretches using the same 20SA-CASSCF(8,8)/cc-pVDZ level of theory. This active space was chosen after careful benchmarking of the vertical excitation energies with different active spaces, which is presented in Appendix Section A. The optimization was carried out using the Gaussian[32]. All of the CASSCF calculations were performed using the COLUMBUS software[33–35].

B. Dynamics

We carried out dynamics starting from each one of the energetically lowest states of CD_2O^{4+}, the first 6 singlet states, and the first 4 triplet states, for a total of 10 independent dynamical runs. Deuterated formaldehyde was chosen for the dynamics because experiments were carried out on the same molecule. The molecular dynamics of the CD_2O^{4+} were calculated in two ways: first, by using only the classical equations of motion for four singly charged atomic ions, and second, using the semi-classical trajectory surface hopping (TSH) approach[19, 20]. For the first approach, because of the highly repulsive nature of tetracation's PESs, it was assumed that the Coulomb explosion of CH_2O^{4+} would lead to the formation of C^+, O^+, D^+, and D^+. Hence, the simulation was performed by treating each charged atom (of +1) as a point charge with a mass equivalent to its atomic mass and a Coulombic force (CF) acting on it due to the other point charges. Alternatively, different partial charges can be used for the Coulombic forces, taking into account that when the tetracation is in equilibrium the charges are not equally distributed in the four atoms. This approach, however, becomes more unrealistic at far distances where all atoms are separated, and they should have a charge of +1. Nevertheless, we compared trajectories using the two different choices of charges, and the results did not change qualitatively. The partial charges on each atom of D_2CO^{4+} were calculated at CASSCF level of theory. A time-step of 0.0082 fs was used, and the simulation was run for 50 fs. 200 initial conditions (ICs) containing the geometries and the initial velocities of CD_2O^{4+} were generated using the harmonic oscillator Wigner distribution of the ground state minimum of CD_2O. The frequencies were calculated at the B3LYP/6-311+G(d) level of theory. The same ICs were used for the dynamics performed with CF as well as TSH. The Wigner distribution code, as implemented in Newton-X, was used[36, 37]. The codes for the dynamics with CF were implemented with Python2.

For the TSH dynamics, the nuclear evolution of the ICs is treated classically using the velocity-Verlet algorithm with a time-step of 0.5 fs. The electronic energies, the non-adiabatic couplings, and the gradients along which these trajectories evolve were calculated at each time-step, on-the-fly at the 8-state averaged (SA8-CASSCF(8,8)/cc-pvDZ) and 6-state averaged SA6-CASSCF(8,8)/cc-pvDZ levels of theories for the dynamics on the singlet and triplet surfaces, respectively. The dynamics were set up to run for a total of 100 fs on S_0 and S_1, although almost all of them were completed by 50 fs. The generation of ICs and the TSH dynamics were done with the help of Newton-X.[36, 37] A trajectory was eliminated if the total energy at any time-step deviated by more than 0.5 eV from the energy at the previous time-step or at time zero. The 'Fewest Switches Surface Hopping' (FSSH) algorithm implemented in Newton-X was used to calculate hopping probability between non-adiabatic surfaces[38]. After a trajectory hops due to non-adiabatic coupling, its momentum is re-scaled along the derivative coupling vector to conserve the total energy. Also, in the case of frustrated hopping, i.e., when a trajectory does not have enough energy to hop, the direction of its momentum is left unaltered. To deal with decoherence, the decoherence correction method of Persico and Grannuci[39] was used with a correction factor of 0.1 Hartree[40]. The results of the TSH dynamics are shown for the first 50 fs, even though they were run for 100 fs. This is because most of the interesting phenomena occurred in the first 50 fs of the dynamics. Also, the trajectories on many states did not survive after 50 fs due to failure in the CASSCF calculations convergence or energy conservation failure (See Appendix Figure 9 where the C-O bond length vs. time is being plotted for 100 fs).

C. Experimental

The experiment makes use of a velocity map imaging (VMI) apparatus [41] outfitted with a Tpx3Cam camera [42]. The laser pulses originate from a commercial amplified Ti:sapphire laser system, which produces 30 fs laser pulses with 1 mJ of energy at a repetition rate of 1 kHz. The pulses are spectrally broadened in a stretched hollow core fiber (S-HCF) [43–46], and compressed using chirped mirrors to a minimum duration of about 6 fs. The pulses (30 fs from the amplifier or 6 fs from the S-HCF) are focused into our VMI apparatus to ionize molecules.

The sample, deuterated formaldehyde, is obtained by sublimation of paraformaldehyde-D_2 (purity 98%, Sigma–Aldrich) at 50–60 °C. A skimmed molecular beam of sample molecules intersects the laser in the VMI apparatus.

The ionized fragments are accelerated toward the microchannel plates (MCP) and phosphor screen under velocity-mapping conditions. The fluorescence from hits on the phosphor screen is imaged onto the camera with an $f/0.95$ lens. The 1 ns precision of the Tpx3Cam is sufficient to resolve the ion momenta along the time-of-

States	Sym	E (eV)	3-body # traj	4-body # traj	failed # traj
S_0	$1\ ^1B_1$	0.00	120	9	71
S_1	$1\ ^1A_1$	0.19	94	17	89
S_2	$2\ ^1A_1$	0.43	71	50	79
S_3	$1\ ^1A_2$	0.90	68	51	81
S_4	$3\ ^1A_1$	2.24	76	62	62
S_5	$1\ ^1B_2$	2.68	34	119	47
T_1	$1\ ^3B_1$	-1.60	137	41	22
T_2	$1\ ^3A_2$	-0.37	65	105	30
T_3	$1\ ^3B_2$	0.24	118	57	25
T_4	$2\ ^3B_1$	1.22	81	90	29

TABLE I. Number of trajectories leading to three body ($CO^{2+}/D^+/D^+$) or four body ($C^+/O^+/D^+/D^+$) dissociation on each electronic state. Also provided are the respective energies and symmetries of electronic states at the Franck-Condon region. The electronic structure calculations were performed at the CASSCF(8,8)/cc-pVDZ level of theory. A total of 200 trajectories were run on each state, and several of them failed before it was clear what the dissociation products were. Number of failed trajectories is shown in the last column.

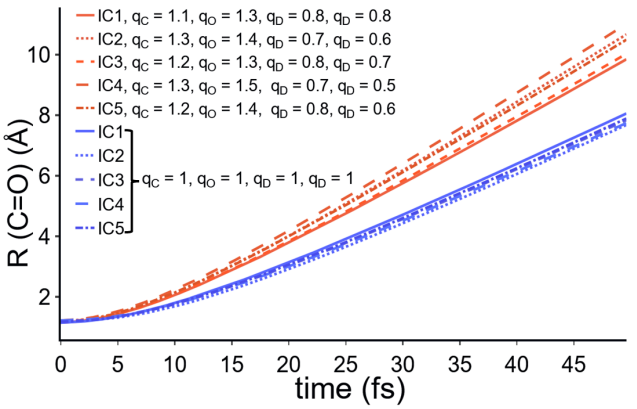

FIG. 1. CO bond length(R(C=O)) plotted for five trajectories (IC1-IC5, starting from different initial conditions (IC)) simulated with CF in two ways, a) q = +1 was assumed on each of the atoms, C, O, D and D, and b) q = Mulliken's partial charges on each of the atoms were calculated at MSCCF(8,8)/cc-pVDZ level of theory. The Mulliken charges for each trajectory are shown in the inset.

flight (ToF) direction, which can be used to reconstruct the full 3D-vector momenta of ions [42, 47].

III. RESULTS AND DISCUSSION

A. Dynamics

We first present the results of TSH dynamics, which were run on 6 singlet and 4 triplet states of CD_2O^{4+}. Table I summarizes the fragmentation patterns observed, showing the number of 3-body ($CO^{2+}/D^+/D^+$) and 4-body ($C^+/O^+/D^+/D^+$) dissociative products formed by running the TSH dynamics on each state. The electronic energies of states at vertical ionization are also shown relative to the lowest singlet state of the tetracation S_0 (note that the lowest energy state is a triplet, T_1). Out of 10 surfaces (6 singlets and 4 triplets) on which TSH dynamics were run, all supported the formation of CO^{2+} to various degrees. The dynamics on electronic states S_0, S_1, T_1, and T_3 yielded a high number of 3-body products. On the other hand, S_5 and T_2 are rich in producing 4-body products. In order to understand better the origin of this behavior, we focus more on the ground state (i.e. T_1 surface) since the dynamics on the other states can be explained analogously.

For the CF approach, there are no excited states, so only one set of trajectories were run. Furthermore, one has to decide what charge should be used on each atom. Two obvious choices exist. One is to use +1 charge on each atom, since this is the correct dissociation limit, after Coulomb explosion when all the atoms are far away from each other. But when the atoms are still close to each other and there is still some bonding interactions, the charges are not +1. In that case, partial charges can be obtained from the ab initio calculations. These charges will be more appropriate initially, but a fractional charge on an atom is unrealistic when atoms are far from each other. We compared the dynamics for a few representative trajectories using the two different sets of charges, and the results are shown in Figure 1. Clearly, in both scenarios, atoms fly away from each other, exhibiting similar physics. So, we will continue our comparisons with a full set of trajectories using +1 charges, which represent better the dissociation limit.

We then compare the TSH results with the dynamics run with CF and point charges. Results for the ground state T_1 are shown here, while results for the other states are shown in the Appendix Section D. The idea is to compare the dynamics run with a semi-classical method (TSH) with entirely classical dynamics and check the validity of this elementary approach. Figure 2 shows the evolution of C-O and two C-D bond lengths with respect to time for the dynamics run on the T_1 state of CD_2O^{4+} for both sets of dynamics. The comparison between the CF and TSH dynamics clearly shows that the dynamics of the dissociation of the two C-D bonds are similar between the two approaches, which means that the two D^+ dissociate fast and classically, but the C-O dissociation behaves very differently. C-O dissociation using CF is very fast. On the other hand, only 41 trajectories out of 200 on T_1 dissociate within 50 fs when using TSH. Even those that dissociate within 50 fs are much slower than in the classical case. This is because classical mechanics cannot capture the bonding nature of CO^{2+} once the two Ds break away quickly. The internal coordinates vs. time are also plotted for the TSH dynamics on other electronic states in Appendix Figure 9. All of the C-O plots show that the dissociation of CD_2O^{4+} can indeed form metastable CO^{2+} through all the electronic states

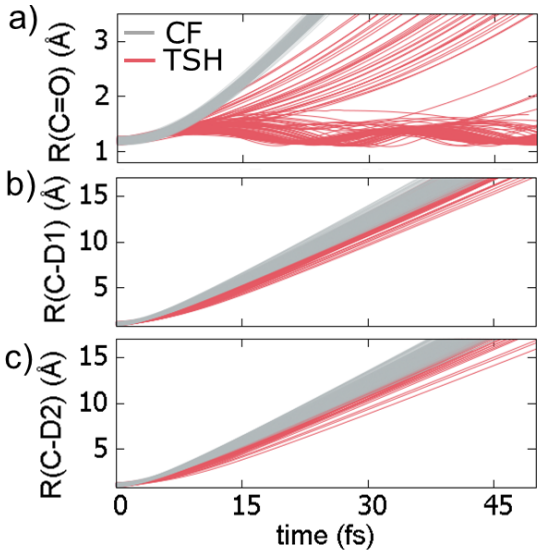

FIG. 2. Bond length of a) C-O, b) C-D1, and c) C-D2 plotted vs time for the dynamics performed on T_1 state of CD_2O^{4+} using simple classical Coulombic forces (CF) and trajectory surface hopping (TSH).

of CD_2O^{4+}.

B. Potential energy surfaces

In order to understand the dynamics better, we examine the potential energy surfaces (PESs) of the electronic states of CD_2O^{4+} along the C-O and C-D (or C-H) bonds. Figure 3 shows the adiabatic T_1 3 dimensional surface of CD_2O^{4+} plotted against the C-O stretch and the symmetric C-D stretch. According to Figure 2 the dissociation dynamics of the C-D bonds are symmetric, i.e., both are stretched at the same time, so it makes sense to plot PESs against the symmetric C-D stretch. The 200 TSH trajectories are also superimposed on Figure 3. One can notice the bifurcation of trajectories in green and red. The 41 trajectories in red represent the trajectories where C-O dissociates to form the product $C^+/O^+/D^+/D^+$, whereas the trajectories shown in green form $CO^{2+}/D^+/D^+$. Some red trajectories overlap with the green trajectories, because they also dissociate but at longer C-D bond lengths, which are out of the range shown in this PES. The shape of the PES makes clear the reason for the formation of CO^{2+}. When the CD bonds are short (in intact CD_2O^{4+}), C-O is directly dissociative. However, as soon as the CD bonds start breaking, a minimum is formed along the CO bond, which can trap the population. Most of the trajectories (green lines in the figure) follow the gradient and get trapped in the well, although there are several trajectories that escape (red lines) and break the CO bond. In order to better understand the dynamics, we now consider cuts of the 3D PES along the C-O stretch for certain values of C-D stretch.

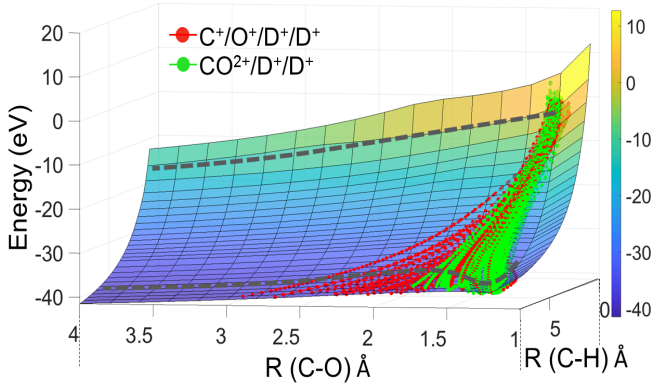

FIG. 3. 3D PES of the T_1 state along C-O and symmmetric C-H bonds. Bifurcation of trajectories into formation of the products $CO^{2+}/D^+/D^+$ (green) and $C^+/O^+/D^+/D^+$ (red) is shown. The 2D cuts of the PES along the grey dashed lines are shown in Figure 4.

Figure 4a shows CD_2O^{4+} PESs along the C-O stretch for 6 singlet and 4 triplet states. The configurations describing these states are shown in the Appendix, Section B. As expected, the PESs of all the states are dissociative because of their Coulombic nature. The same is true for the PESs along the C-D stretch, as can be seen in Figure 5. One can hypothesize (and it is confirmed by the dynamics) that since the mass of oxygen is 16 times (or 8 times) heavier than hydrogen (or Deuterium), the C-D dissociation should be very fast compared to C-O dissociation in CH_2O^{4+} (or CD_2O^{4+}). However, in that scenario, once the D's dissociate away from CD_2O^{4+}, the dissociation of C-O then should be controlled by the PESs of CO^{2+}. Figure 4b shows the PESs of CO^{2+}, calculated using SA20-CASSCF(8,8)/cc-pVDZ, which demonstrate that a few electronic states with high energy barriers can support the formation of metastable CO^{2+}. Some of these states are $1\ ^1\Sigma^+$, $1\ ^1\Pi$, $1\ ^3\Pi$, and $1\ ^3\Sigma^+$ with energy barriers of 3.2 eV, 2.5 eV, 1.3 eV, and 2.4 eV along the C-O stretch. Please refer to Table II to correlate CH_2O^{4+} states to the CO^{2+} states, and for excitation energies of CO^{2+}. The deep or shallow wells along the C-O stretch can hence support the formation of CO^{2+}. The depth of these wells defines the lifetimes of the metastable CO^{2+}, determining the tunneling rate through barriers.

TSH dynamics, in general, cannot capture the tunneling of metastable populations through the energy barriers since it is a semi-classical method. However, the lifetimes of metastable CO^{2+} have been discussed previously in the literature. Both experimental [48–51] and theoretical results [52] have shown that the lifetimes of the first few vibronic states can range from a few picoseconds to seconds for states such as the ν=0,1 vibrational states of $1^3\Pi$. Since the ToF of the VMI apparatus used in the experiment is on the order of μs (which is more than the lifetimes of most of the states), the experiments

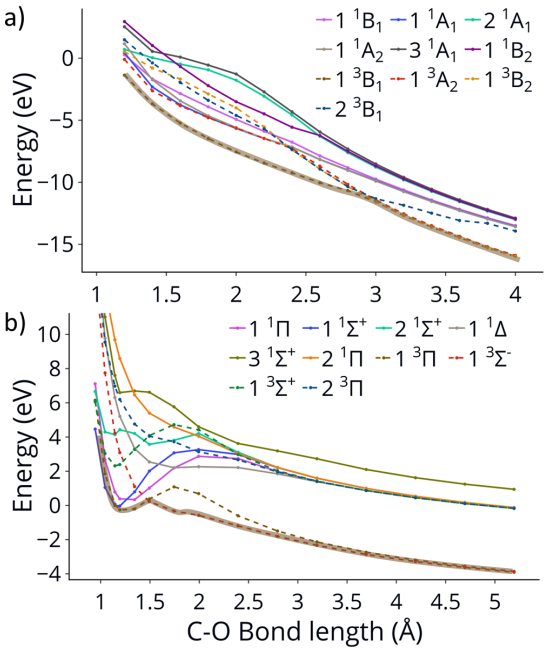

FIG. 4. PESs of 6 singlets (solid line) and 4 triplets (dashed line) calculated along the C-O stretch for a) CH_2O^{4+}, and b) CO^{2+}. The calculations were carried out at CASSCF(8,8)/cc-pVDZ level of theory. All of the energies in a) are with respect to the energy of the lowest singlet state at the neutral equilibrium geometry of CH_2O^{4+}, and in b) are with respect to the energy of the CO^{2+} ground state minimum. Here, 1 $^3B_1$ and 1 $^3B_2$ of D_2CO^{4+} converts into 1 $^3\Pi$ state of CO^{2+}. For all the correlations between D_2CO^{4+} and CO^{2+} states, see Table II. The adiabatic T_1 state is highlighted with a thick transparent brown line.

D_2CO^{4+} states	Corresponding CO^{2+} states	VEs (eV) of CO^{2+} states
1 $^3B_1$ + 1 $^3B_2$	1 $^3\Pi$	0.08
2 $^3B_1$ + 2 $^3B_2$	2 $^3\Pi$	6.97
1 $^3A_2$	1 $^3\Sigma^-$	4.21
1 $^3A_1$	1 $^3\Sigma^+$	2.31
1 $^1B_1$ + 1 $^1B_2$	1 $^1\Pi$	0.79
1 $^1A_1$	1 $^1\Sigma^+$	0.00
2 $^1A_1$	2 $^1\Sigma^+$	4.14
2 $^1A_2$	1 $^1\Sigma^-$	9.81
3 $^1A_1$ + 1 $^1A_2$	1 $^1\Delta$	6.31

TABLE II. Correspondance of D_2CO^{4+} states to CO^{2+} states. The corresponding vertical excitation energies (VEs) of CO^{2+} are also provided with respect to the S_0 minimum of CO^{2+} which is $^1\Sigma^+$.

can capture the signature of metastable CO^{2+}.

C. Comparison with the experiment

We compare our CF and TSH calculations with the experiment using a native frame analysis in order to account for the sequential dissociation of CO^{2+}. The experimental signal has been presented in detail in Ref.[21] and here the relevant plot is used for comparisons with theory. Native frame analysis has been very helpful in distinguishing between the sequential vs. concerted dynamics using the Jacobi coordinates, especially in the case of three body dissociation[7, 53]. This analysis aims to compare an observable, here, the kinetic energy of CO^{2+} with respect to the momenta of the two D^+ (KE_{CO}) calculated using theory and experiment in the native frame. Note that due to complications in the measurements, CH_2O^{4+} was used rather than CD_2O^{4+} for the native frame analysis. The relevant axes for the analysis can be seen in Figure 6e, where the x-axis is along the sum of momenta

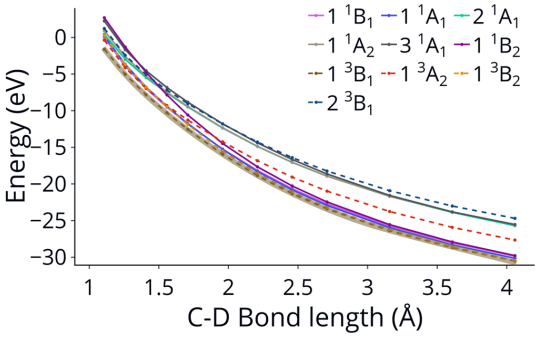

FIG. 5. PESs of 6 singlets (solid line) and 4 triplets (dashed line) calculated along the C-D (or C-H) stretch for CD_2O^{4+} (or CH_2O^{4+}). The calculations were carried out at CASSCF(8,8)/cc-pVDZ level of theory. All of the energies are with respect to the S_0 state at the neutral equilibrium geometry of CH_2O^{4+}. The adiabatic T_1 state is highlighted with a thick transparent brown line.

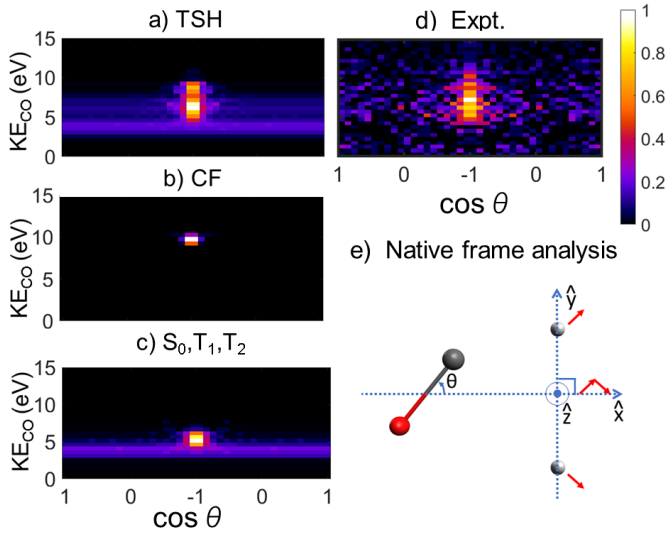

FIG. 6. Distributions of dissociation energies of CO^{2+} (KE_{CO}) against cos of the angle defined in the native frame θ for a) TSH trajectories on all the 10 states, b) CF, c) TSH trajectories on T_1, T_2, and S_0, and d) Experimental results for a 6 fs laser pulse (Expt.) (taken from Ref. [21]). e) Figure explaining the axes used for the native frame analysis.

vectors of both the D cations (red arrows), the y-axis is along the difference between them, and the z-axis is the cross vector of x and y axes. Also, θ is the angle of rotated CO^{2+} with respect to the x-axis. In order to calculate KE_{CO} when running TSH we need to switch to classical Coulombic forces after the trajectories fail because they almost never reach a point where the Coulomb interactions are zero. At the point of switching, there is an abrupt change from a PES calculated using quantum chemistry (QPES) and CF (CPES), as seen in Appendix Figure 8. Hence, the KE_{CO} values were shifted with the difference between the QPES and CPES at the last point of the TSH trajectory, such that the abrupt change between QPES to CPES is compensated. This correction is described in detail in Appendix Section C.

In Figure 6, we compare the experimental results obtained with a 6 fs laser to our TSH and CF calculations analyzed in the native frame. The figure shows the 2D distribution of calculated KE_{CO} along the cos θ values for a) TSH dynamics on all states, b) CF dynamics, c) TSH dynamics on T_1, T_2, and S_0, and d) experiment. Clearly, the CF plot does not resemble the experimental distribution of KE_{CO}, while the experimental (expt.) and TSH results are qualitatively similar. For example, the maximum dissociation energies of CO^{2+} (i.e., KE_{CO}) are mostly concentrated around 7-8 eV in both TSH and expt. And, the KE_{CO} values are mostly concentrated at $\cos\theta$ value of -1.

It is interesting to understand the physical origins of the different features in the TSH plot. The low energy but broad angular distribution arises from rotation of the CO^{2+} after both D^+ ions break off. Given the lifetime of CO^{2+}, its formation in some TSH trajectories left it with some angular momentum before dissociation. This broad distribution is not as prominent or clear in the experimental data (Figure 6d). This could be because i) the noise in experiments hides them, ii) the rotational time scale is comparable to the dissociation time scale of CO^{2+} (i.e., a few ps), iii) H_2CO^{4+} may form less metastable CO^{2+} as compared to D_2CO^{4+}.

The broadness of the KE_{CO} distribution from the TSH calculations arises from the fact that the tetracation population was formed on multiple electronic states (here, 6 singlets and 4 triplets were considered in TSH dynamics). This is clear by comparing Figure 6a to Figure 6c, where a similar plot has been plotted for the dynamics on the 3 lowest states, i.e., T_1, T_2, and S_0, where the KE_{CO} distribution is much narrower. One can also notice that there are some differences in the range of KE_{CO} distribution in the experiment and TSH calculations. There is no signal after $\sim$10 eV in the latter, while in the experiment the energy extends to somewhat higher energies. This could happen because only 10 states were considered for the TSH dynamics. The inclusion of more electronic states in TSH can make the KE_{CO} distribution broader, bringing it into closer agreement with the experiment. Another reason for discrepancies between experiments and TSH results can be that all the states were equally populated with 200 initial trajectories to do the dynamics. Knowing the ionization probability to ionize CD_2O to a certain state of the tetracation can improve the TSH results. However, it is very difficult to guess the probability during strong-field ionization due to the complications that can arise during such processes[54]. Hence, the simplest way to analyze the calculations was to assume that all the states were equally populated and vertically ionized.

D. Conclusion

In summary, in this work, a comparison between theoretical calculations and experiment demonstrated that a simple classical description of the dynamics of Coulomb explosion using classical forces and the assumption that a single charge exists on each atom is not adequate. The presence of metastable fragments requires a quantum mechanical description of the bonding between atoms. Furthermore, the results also indicate that ionization to multiple electronic states occurs. These results provide an important insight on how to properly model Coulomb explosion imaging, in order to properly utilize the technique to obtain structures of molecules.

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Appendix A: Vertical energies benchmarked with different active spaces of CASSCF

Presented in Table III are the vertical energies of H_2CO^{4+} (or D_2CO^{4+}) calculated with different active spaces of CASSCF. It can be seen that all levels of theory provide similar energies. However, more states were averaged than the states involved in the dynamics because higher states can get involved in the dynamics at later time scales. For example, 8 singlet states were averaged to calculate the electronic energies of 6 singlets that were involved in the TSH dynamics. Similarly, 6 triplet states were averaged to do the TSH dynamics with 4 triplets. Note that the quintets are higher in energy and, hence, were not considered for the dynamics.

Singlets	S_0	S_1	S_2	S_3	S_4	S_5
C_{2v} Symmetry	$^1B_1$	$^1A_1$	$^1A_1$	$^1A_2$	$^1A_1$	$^1B_2$
5SA-CAS(6,6)/cc-pvdz	0.00	0.22	0.59	0.93	2.51	-
5SA-CAS(6,8)/cc-pvdz	0.00	0.17	0.82	0.83	2.55	-
8SA-CAS(8,8)/cc-pvdz[a]	0.00	0.29	0.52	0.93	2.15	2.71
20SA-CAS(8,8)/cc-pvdz[b]	0.00	0.19	0.43	0.90	2.24	2.68
Triplets	T_1	T_2	T_3	T_4		
C_{2v} Symmetry	$^3B_1$	$^3A_2$	$^3B_2$	$^3B_1$		
4SA-CAS(6,6)/cc-pvdz	-1.72	-0.26	0.21	1.38		
6SA-CAS(8,8)/cc-pvdz[c]	-1.52	-0.33	0.22	1.25		
20SA-CAS(8,8)/cc-pvdz[d]	-1.60	-0.37	0.24	1.22		
Quintets	Q_1	Q_2	Q_3	Q_4		
C_{2v} Symmetry	$^5B_1$	$^5A_1$	$^5A_2$	$^5A_1$		
4SA-CAS(8,8)/cc-pvdz	1.53	3.69	4.01	4.48		

[a] used for TSH dynamics
[b] used for PES calculations
[c] used for TSH dynamics
[d] used for PES calculations

TABLE III. Energies of the tetracation states at the neutral geometry of formaldehyde calculated with different active spaces of CASSCF. All the energies at the particular level of theory are with respect to the S_0 energy at the neutral geometry.

Appendix B: Electronic configurations of states of H_2CO^{4+}

The electronic configurations of 6 singlets and 4 triplets of H_2CO^{4+} at the neutral geometry are provided in Figure 7. The electronic configuration of H_2CO is also provided for reference. This can help readers identify the symmetry as well as the character of the states provided the holes in the orbitals. For example, S_0 state has a hole in $5a_1$ and $1b_1$ orbitals. The tensor product of a_1 and b_1 gives the symmetry of S_0 state, which is nothing but b_1.

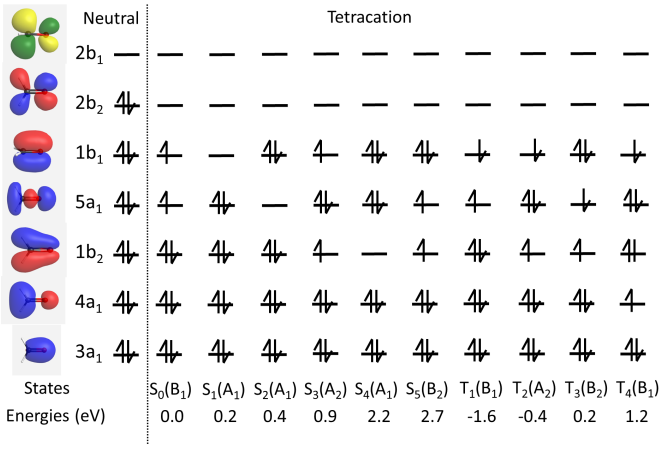

FIG. 7. Electronic configuration of first 6 singlets and 4 triplets at the neutral geometry of CH_2O^{4+}. The electronic configuration of the ground state (S_0) of CH_2O is also provided, separated by a dashed line on left-hand side.

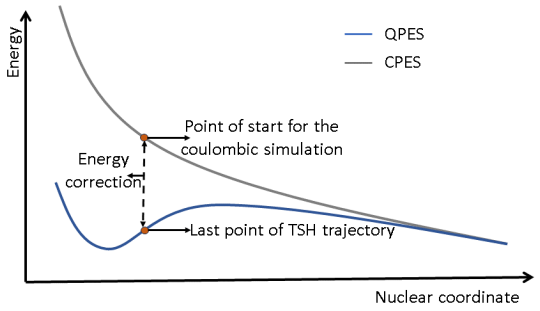

FIG. 8. Cartoon explaining the correction in energy shift made to KE_{CO}.

Appendix C: Native frame analysis

The main details about the analysis have been provided in the paper. As said in the paper, for the theoretical analysis, the momenta of two Ds, C and O, were taken from the last step of the TSH trajectories and then simulated with CF in order to get the final momenta of the four bodies when they are far away, and there are no interactions between them. However, there is an abrupt change from a PES calculated using quantum chemistry (QPES) and CF (CPES), as seen in Figure 8. Hence, the KE_{CO} values were shifted down with the difference between the QPES and CPES at the last point of the TSH trajectory (shown with the double-headed dashed arrow in Figure 8). The computed data were then distributed into 2D bins of KE_{CO} and cos θ with bin sizes of 0.5 eV and 0.1, respectively. Also, in the case of TSH trajectories that ended with CO^{2+}, the value in each bin was distributed equally to all the cos θ bins by dividing it by the total number of bins (which equals 20 here). The idea of equally spreading the cos θ value among all the bins is to conserve the angular momentum of rotating CO^{2+} among all the θ values. The final distribution is normalized w.r.t. the maximum KE_{CO} value and mirrored around the -1 value of cos θ.

Appendix D: TSH dynamics on other electronic states of D_2CO^{4+}

Figure 9 shows the evolution of C-O bond length vs time for the dynamics ran on different electronic states. It can be seen that the metastable CO_2^+ are being formed on each electronic state.

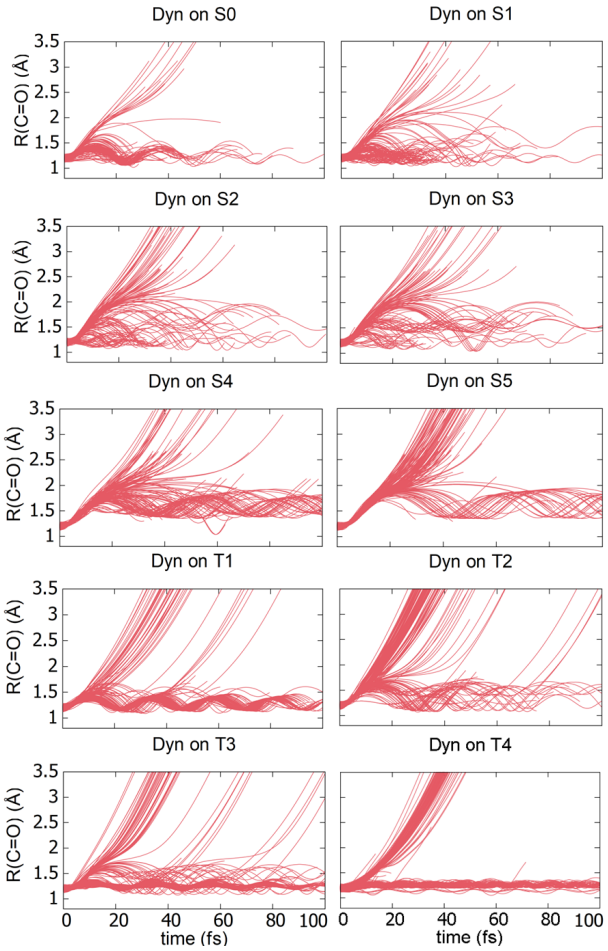

FIG. 9. C-O vs time plotted for TSH dynamics (dyn) ran on different electronic states.

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