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An axis-specific rotational rainbow in the direct scatter of formaldehyde from Au(111) and its influence on trapping probability†

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The conversion of translational to rotational motion often plays a major role in the trapping of small molecules at surfaces, a crucial first step for a wide variety chemical processes that occur at gas–surface interfaces. However, to date most quantum-state resolved surface scattering experiments have been performed on diatomic molecules, and little detailed information is available about how the structure of nonlinear polyatomic molecules influences the mechanisms for energy exchange with surfaces. In the current work, we employ a new rotationally resolved 1 + 1' resonance-enhanced multiphoton ionization (REMPI) scheme to measure the rotational distribution in formaldehyde molecules directly scattered from the Au(111) surface at incidence kinetic energies in the range 0.3–1.2 eV. The results indicate a pronounced propensity to excite *a*-axis rotation (twirling) rather than *b*- or *c*-axis rotation (tumbling or cartwheeling), and are consistent with a rotational rainbow scattering model. Classical trajectory calculations suggest that the effect arises—to zeroth order—from the three-dimensional shape of the molecule (steric effects). Analysis suggests that the high degree of rotational excitation has a substantial influence on the trapping probability of formaldehyde at incidence translational energies above 0.5 eV.

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1. Introduction

The interconversion of translational and rotational energy in molecule–surface collisions can play a key role in the mechanism for adsorption,^{1–5} which is the crucial first step for a large number of chemical processes that take place at gas–surface interfaces.^{6,7} This has motivated a large number of quantum-state resolved measurements of rotational distributions in molecules scattered or desorbed from surfaces, which provide detailed information not only on the mechanism for trapping/desorption,^{2,8–10} but also on orientation dependence of molecule–surface interactions (steric effects).^{11–16} The rotational distributions obtained from direct (*i.e.* single-collision-event) molecular scattering experiments are often non-thermal, and striking features can arise from so-called “rotational rainbow” effects, which occur when there is a singularity in the classical rotational

energy distribution due to an extremum in $J(\gamma)$, (where J is the rotational angular momentum in the scattering product and γ is the orientation of the incoming molecule).¹⁷ Such effects are analogous to those caused by extrema in the classically achievable scattering angle of light from water droplets—hence the name. The consequence of an extremum in $J(\gamma)$ is that there is a range of initial γ leading to approximately the same J , such that population in the scattering products accumulates at that J . Quantum effects lead to broadened maxima, rather than sharp singularities, in the rotational state distributions observed in molecular beam scattering experiments.¹⁸ Classical and quantum calculations based on model potentials have succeeded in reproducing the key features of the experimentally observed rotational rainbow effects in a variety of systems, displaying diverse behavior.^{19–25}

The excitation of molecular degrees of freedom becomes increasingly important in the dissipation of incident kinetic energy during molecule/surface collisions as the number of atoms in the molecule is increased. In fact, rotational excitation has been invoked as a key component of the trapping mechanism in a number of polyatomic molecule–surface systems.^{4,5} However, to date, almost all experiments that probe quantum-state resolved rotational excitation at surfaces have been limited to diatomic or linear polyatomic^{26,27} molecules. The degree to which nonlinear polyatomic molecules are rotationally excited

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during collisions with surfaces remains an experimentally unexplored area. We are aware of only two previous surface scattering experiments that have probed the rotational distribution in a scattered nonlinear polyatomic molecule: CH₄ from LiF(100)²⁸ and NH₃ from Au(111).^{25,29} The CH₄ study, which was conducted at an incidence energy of 0.075 eV and an incidence angle of 60°, found no significant deviation from a thermal rotational distribution. On the other hand, the NH₃ study, which was conducted at an incidence energy of 0.24 eV and an incidence angle of 45°, found a strong propensity for a rotationally cold NH₃ beam to scatter elastically into low-*J* states. A propensity to excite low-*K* states (tumbling rather than twirling) was also observed. The results were interpreted in terms of a dynamical steering effect, which orients the symmetry axis of the NH₃ molecule along the surface normal prior to collision.

In the current work, we report rotationally-resolved molecular beam scattering of formaldehyde from Au(111). We report an axis-specific rotational rainbow in the direct scattering channel. Unlike in the case of NH₃, direct scatter of formaldehyde from Au(111) leads to a clear propensity for excitation of rotational states with a high degree of rotational energy about the symmetry axis. The results are interpreted with the aid of classical trajectory calculations. To our knowledge, the present work is the first report of rotationally resolved surface scattering of an asymmetric top molecule and the first report of a high-*J* rotational rainbow in a nonlinear molecule. The work demonstrates the ability to probe the highly structured steric effects that govern the interactions between polyatomic molecules and surfaces, and the results suggest that rotational excitation contributes to the trapping probability of formaldehyde on Au(111) at high incidence kinetic energies.

II. Methods

A. Experimental

The essential details of the experiment are described in ref. 30. A pulsed beam of formaldehyde was generated by cracking solid paraformaldehyde (97% purity) in a sample holder inside the nozzle of a home-built pulsed solenoid valve, described in ref. 31. The tip of the nozzle was heated to a temperature of 104–130 °C and the sample holder was maintained at a cracking temperature of 60–80 °C. The molecular expansion is sent through a 2 mm electroformed skimmer (Ni Model 2, Beam Dynamics, Inc.) located 6 cm from the nozzle and through a second 2 mm aperture, located 20 cm downstream, before entering the ultra-high vacuum (UHV) chamber, where the surface scattering experiments are performed. We calculate the peak density of the incoming beam in the UHV chamber to be $7 \times 10^{11} \text{ cm}^{-3}$, so that the mean free path of scattered molecules traveling through the incoming beam can be conservatively estimated to be $\gtrsim 1 \text{ m}$, and collisions between molecules can be excluded.

The incidence kinetic energy (E_i) of the incoming beam of formaldehyde was varied over the range 0.1–1.2 eV, by using different backing gas mixtures (see Table SI of the ESI†).

The stagnation pressure behind the nozzle was also varied over the range 3–12 bar to achieve a 3–15 K rotational temperature (T_{rot}) in the incoming molecular beam. Under these conditions, the incoming beams used in this study had no significant population in levels with $K_a > 1$ or $J > 4$.

The incoming beam was scattered at near-normal incidence angle from a Au(111) crystal (Mateck). Before each experiment, the Au(111) crystal is cleaned by sputtering with an Ar-ion gun (LK Technologies NGI3000) and annealed for 20 minutes at 900 K. Surface cleanliness is verified by Auger electron spectroscopy (Staib Instruments, Inc. ESA-150). The temperature (T_s) of the Au(111) crystal was monitored by a thermocouple (K type), and could be cooled by liquid nitrogen or heated resistively to achieve temperatures in the range 100–1000 K.

The rotational distribution in the vibronic ground state was monitored in the incoming and scattered molecular beam *via* a $1 + 1'$ REMPI scheme, which we recently reported.³² Briefly, the frequency-doubled output of a pulsed dye laser (Sirah Cobra Stretch SL, operating with LDS 698 or 722 dye), pumped at 10 Hz by a Nd:YAG laser (Continuum Powerlite 8010), was scanned across the rotationally resolved $\tilde{A}^1A_2 \leftarrow \tilde{X}^1A_1 (4_0^1)$ transition at 353 nm. (We use the notation $4_{v_4''}^{v_4'}$ to denote the number of quanta of out-of-plane wagging vibration ν_4 in the upper (v_4') and lower (v_4'') electronic state, respectively.) The pulse energy was typically 5–7 mJ and the beam was collimated to a diameter of $\sim 3 \text{ mm}$. A small portion of the dye laser fundamental was coupled into a high-precision wavemeter (HighFinesse, WS7) for frequency calibration, to an accuracy of $\pm 0.02 \text{ cm}^{-1}$.³¹ Variation in the laser intensity over the scan was monitored by recording the shot-by-shot pulse energy behind the chamber. After a delay of 20 ns, molecules were ionized from the \tilde{A} state by a single vacuum ultraviolet (VUV) photon at 157 nm, obtained from a molecular F₂ laser (GAM EX350). The VUV beam (2–8 mJ pulse energy) was counter-propagated relative to the dye laser beam and was focused into the interaction region by a $f = 58.1 \text{ cm}$ CaF₂ lens. The experiment is operated in a regime where the signal is linear with respect to both laser pulses, and the relative integrated ion intensities are directly proportional to the $\tilde{A} \leftarrow \tilde{X}$ absorption cross section to within $\sim 20\%$ uncertainty.³² Molecular cations were collected by a set of ion optics and detected on a 2-stage microchannel plate assembly (Hamamatsu F1552) in chevron configuration.

At $E_i = 0.1 \text{ eV}$, the scattering is dominated by a trapping/desorption mechanism in which molecules physisorb to the surface before undergoing thermal desorption. At higher incidence energies ($E_i = 0.3\text{--}1.2 \text{ eV}$), we observe distinct channels for trapping/desorption and direct (single-impact) scattering. These channels have been characterized in detail in ref. 30. At incidence energies between 0.33–0.39 eV we probe the scattered molecules at a distance of 9 mm from the surface, 3 mm above the incoming beam. This allows us to capture a narrow range of near-specular scattering angles, in which the direct scatter channel dominates. At incidence energies greater than 0.39 eV the direct scatter channel dominates over a wider range of scattering angle. At high E_i , the direct scattering channel can



also be resolved from the trapping/desorption channel and from the incoming beam *via* ion time-of-flight (TOF) to the detector.³⁰ The scattering products in the $E_i = 0.48$ – 1.2 eV range were thus monitored 3 mm from the surface, such that a wide range of scattering angles is probed. No significant variation of rotational distribution was found over the range of distances and angles probed in our experiment.

B. Fitting and simulation

For comparison with our measured spectra, we calculate the frequency and absorption intensity of each line using a Watson asymmetric top Hamiltonian, with the parameters of ref. 33 for the \tilde{A}^1 state and the parameters of ref. 34 for the ground state. The lines in the b -type $\tilde{A} \leftarrow \tilde{X} (4_0^1)$ transition are sufficiently dense that few individual rotational lines are fully resolved in the spectra of rotationally hot scattering products, obtained with a dye laser resolution of ~ 0.1 cm⁻¹. Therefore, a Boltzmann plot analysis based on individual integrated linestrengths is not possible. However, our resolution is sufficient to obtain many partially resolved lineshapes that give rise to sharp features in the spectra, even in the hottest rotational distributions observed. We have made detailed manual feature-by-feature analyses of the rotational structure, and we have supplemented this analysis by fitting the observed spectra to simulated spectra obtained by modeling the rotational distribution. Additional details about the fitting method are given in Section SII of the ESI.[†]

III. Results and analysis

A. Trapping/desorption channel: dependence of the rotational distribution on T_s

At the lowest incidence kinetic energy employed in our study ($E_i = 0.11$ eV), the trapping/desorption mechanism dominates.³⁰ We have recorded REMPI spectra of molecules scattered in this regime for surface temperatures ranging from 140–480 K. An analysis of the spectra (see Section SIII of the ESI[†]) reveals no significant departure from a thermal Boltzmann distribution of rotational state population and indicates nearly complete equilibration of rotational degrees of freedom to the surface temperature.

B. Direct scatter channel: dependence of the rotational distribution on E_i

The $\tilde{A} \leftarrow \tilde{X} (4_0^1)$ spectra of formaldehyde molecules, scattered from Au(111) at incidence kinetic energies over the range 0.3–1.2 eV and probed at angles where the direct scatter channel dominates, are shown in Fig. 1. The scattered molecules are rotationally much hotter than T_s and the degree of rotational excitation increases as E_i is increased. Even from the “bird’s eye” view presented in Fig. 1, a few key trends are evident in the spectra. First, as E_i is increased, spectral intensity spreads away from the band center at ~ 28300 cm⁻¹, and there is a general shift of intensity towards the lower frequency side of the band because the highest- J progressions extend far to the red of the band center. Although the structure of the hot,

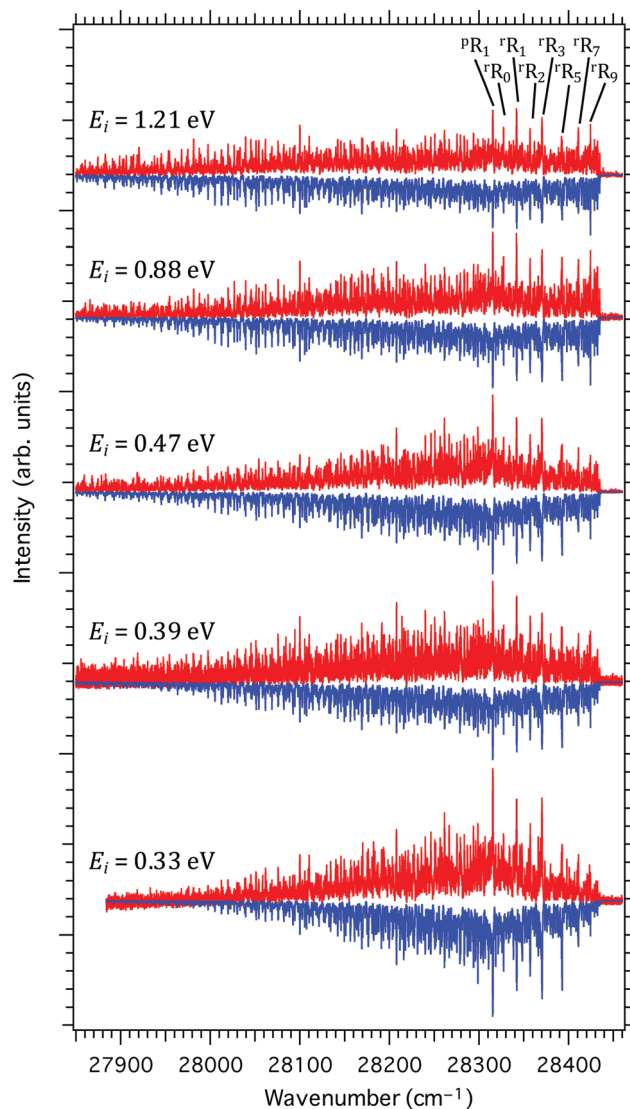


Fig. 1 The $1 + 1'$ REMPI spectrum of the $\tilde{A} \leftarrow \tilde{X} (4_0^1)$ band of formaldehyde molecules, scattered from a Au(111) surface at $T_s = 300$ K, is shown as a function of incidence kinetic energy, E_i (red, upward directed peaks). Shown for comparison are the best-fit simulated spectra obtained using the a -axis Rainbow model (eqn (1)), which will be introduced in Section III B.2. The locations of prominent R-bandheads are labeled at the top of the figure.

asymmetric top spectra is complicated, there is a regular progression of R-bandheads that remains prominent even at the highest E_i , labeled at the top of Fig. 1. (Throughout this work, we use the simplified notation $\Delta K_a \Delta J_{K_a''}$ to label branches of the spectrum. In cases where asymmetry splitting is resolved, we will use the more detailed notation, $\Delta K_a \Delta K_c \Delta J_{K_a'', K_c''} (J'')$ to label individual lines.) At the lowest E_i , most of the intensity is concentrated in the $K_a'' = 0$ – 3 bandheads. However, as E_i is increased, the relative intensity of the $K_a'' = 5$ – 9 bandheads gradually grows in until the progression of bandhead intensities for $K_a = 0$ – 9 becomes almost flat (after taking into account the 3:1 *ortho:para* intensity alternation) at $E_i \geq 0.88$ eV.



1. Comparison with a Boltzmann rotational distribution.

Our first approach to analyzing the spectra was to compare the intensities with those predicted by a thermal Boltzmann population distribution of rotational states. However, a detailed comparison between the observed and calculated intensities reveals large, systematic discrepancies that increase with E_i . The most easily recognized discrepancy is apparent in the high- K_a'' ${}^{\text{r}}R_{K_a''}$ bandhead region at the blue edge of the band.

Fig. 2 compares the observed $1 + 1'$ REMPI spectrum in this region—obtained from molecules scattered at an incidence kinetic energy of 1.21 eV—with the spectrum simulated from the best-fit Boltzmann temperature ($T_{\text{rot}} = 1082$ K). Also shown at the bottom of Fig. 2 is a spectrum simulated using a different model for the population distribution, which we will introduce in Section III B2. The intensities obtained from the Boltzmann distribution are in very good agreement with experiment for the ${}^{\text{r}}R_5$ bandhead at 28393 cm^{-1} (integrated intensity agrees to within 3%). However, the relative intensity of the prominent ${}^{\text{r}}R_7$ bandhead at 28411 cm^{-1} and the ${}^{\text{r}}R_9$ origin at 28424 cm^{-1} are underestimated by the Boltzmann fit by a factor of 1.73 and 2.17, respectively. At higher K_a'' , the discrepancy becomes worse. The intense features in the observed spectrum at $28429\text{--}28435\text{ cm}^{-1}$ —which arise primarily from the prominent ${}^{\text{r}}R_{11}$ progression, as well as from the origins of the ${}^{\text{r}}R_{12\text{--}14}$ progressions—are underestimated by the Boltzmann fit by an integrated factor of 3.4. We can be quite confident about the assignments in this region because the ${}^{\text{r}}R_{K_a''}$ bandheads curve around to the red after $K_a'' = 13$, so that this region is relatively sparse and can only contain lines from the ${}^{\text{r}}R_{10\text{--}15}$ progressions. In order to reproduce the experimentally observed relative intensities in this region, it is necessary to use a Boltzmann rotational temperature of >3000 K. However, the simulated band shape at this temperature exhibits wide overall disagreement with the rest of the observed spectrum. The enhanced intensity in the high- K_a'' ${}^{\text{r}}R_{K_a''}$ bandhead region (relative to that

predicted for a Boltzmann distribution) is reproducible across a wide range of incidence kinetic energies and increases monotonically with E_i .

The low-frequency tail of the band below 28200 cm^{-1} , which contains intensity primarily from ${}^{\text{p}}P_{K_a''}$ branches, is another relatively sparse region that permits a detailed line-by-line analysis of the relative intensities. In Fig. 3, the observed spectrum in this region obtained from a scattered beam ($E_i = 1.21\text{ eV}$) is compared with the simulated spectrum from the best-fit Boltzmann distribution. Again, a simulation from a second model, which we will introduce in Section III B2, is also included in the figure. The observed spectrum contains prominent progressions that can be assigned to the ${}^{\text{p}}P_9$ and ${}^{\text{p}}P_{11}$ branches, labeled in the figure. However, the Boltzmann fit moderately underestimates the contribution of ${}^{\text{p}}P_9$ lines—particularly at low J'' —and grossly underestimates the contribution of ${}^{\text{p}}P_{11}$. For example, the spectrum displays significantly more intensity than the Boltzmann fit at 28110.6 cm^{-1} where the ${}^{\text{p}}P_9(9)$ line is predicted. The features observed at 28076.0 cm^{-1} and 28100.1 cm^{-1} are significantly blue-shifted (by 0.25 and 0.1 cm^{-1} , respectively) in the Boltzmann simulation, which is consistent with missing intensity from the ${}^{\text{p}}P_9(15)$ and ${}^{\text{p}}P_9(11)$ lines. Finally, the relative intensity of the features assignable to ${}^{\text{p}}P_{11}(11\text{--}16)$ is strikingly absent from the Boltzmann fit.

There are other features that appear with high relative intensity in the Boltzmann fit, but are not observed in the spectrum. Among the most prominent of these features are the lines arising from ${}^{\text{p}}P_1(32\text{--}38)$, which are labeled in Fig. 3. The intensity of the features that contain contributions from ${}^{\text{p}}P_1(33\text{--}35)$ at 28082.1 , 28092.6 , and 28102.8 cm^{-1} are notably stronger in the Boltzmann fit than in the observed spectrum. Also, the Boltzmann fit predicts intensity at 28049.7 cm^{-1} and 28101.3 cm^{-1} that arises from the ${}^{\text{p}}P_1(38)$ and ${}^{\text{p}}P_{3,30}(33)$ lines, respectively, whereas there is almost no intensity in the observed spectrum at these frequencies.

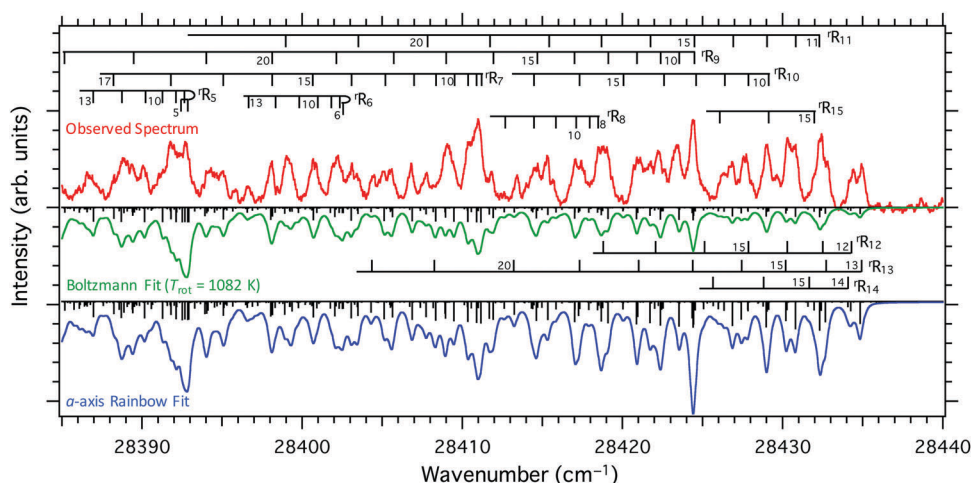


Fig. 2 The high- K_a'' ${}^{\text{r}}R_{K_a''}$ bandhead region of the $\tilde{A} \leftarrow \tilde{X}(4_0^+)$ band of formaldehyde molecules, directly scattered from a Au(111) surface at $T_s = 300\text{ K}$ and $E_i = 1.21\text{ eV}$ is shown (top spectrum, red, upward directed peaks). Shown for comparison are the best-fit simulated spectra obtained from a Boltzmann model (middle spectrum, green, downward directed peaks) and from the α -axis rainbow model (eqn (1)), which will be introduced in Section III B2. The ${}^{\text{r}}R_{5\text{--}15}$ branch transitions are labeled.



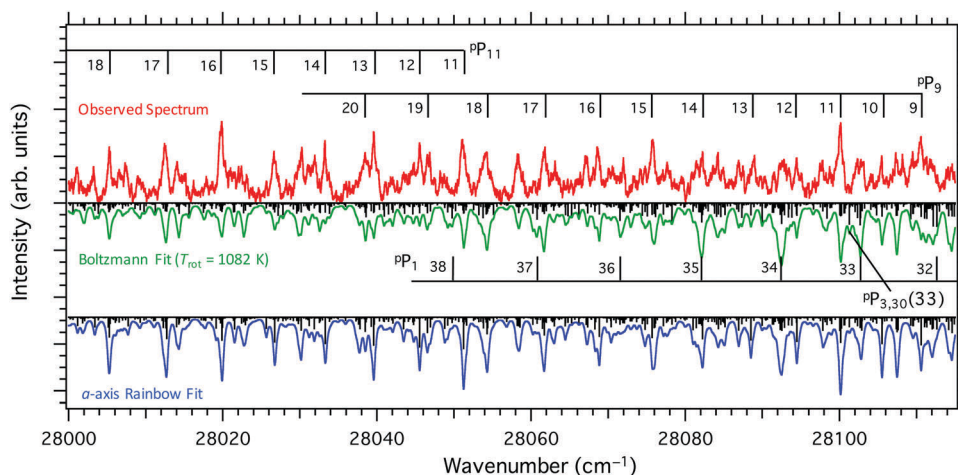


Fig. 3 A portion of the P-P region of the REMPI spectrum of the $\tilde{A} \leftarrow \tilde{X} (4_0^1)$ band of formaldehyde molecules, directly scattered from a Au(111) surface at $T_s = 300$ K and $E_i = 1.21$ eV is shown (top spectrum, red, upward directed peaks). Shown for comparison are the best-fit simulated spectra obtained from a Boltzmann model (middle spectrum, green, downward directed peaks) and from the a -axis rainbow model (eqn (1)), which will be introduced in Section III B2. Selected branches and transitions are labeled.

2. a -axis rotational rainbow model. The lower states involved in the series of transitions labeled in Fig. 3 are similar in total rotational energy. (The $J''_{K_a} = 11_{11} - 18_{11}$ series has $1141 - 1395$ cm^{-1} of rotational energy, whereas the $J''_{K_a} = 32_1 - 38_1$ series has $1231 - 1719$ cm^{-1} of rotational energy.) However, the *distribution* of rotational energy about the distinct internal axes of rotation is very different in these two series. In a near prolate top molecule, K_a is an approximately good quantum number that accurately describes the projection of angular momentum about the a -axis (the CO bond axis). The low- J members of the $K_a = 11$ manifold have almost all of their angular momentum about the a -axis (“twirling” motion), whereas the high- J levels of the $K_a = 1$ manifold have almost all of their angular momentum distributed about the b - and c -axes (“tumbling” or “cartwheeling” motion).

The analysis presented in Section III B1 suggests that the rotational population distribution in the scattered molecular beam deviates from a thermal distribution in a way that favors “twirling” motion with K_a in the range 7–13, over “tumbling/cartwheeling” motion with the same amount of energy. Deviations from a Boltzmann rotational distribution appear quite commonly in experiments on direct scattering of diatomic molecules from surfaces.^{2,9,17} Such effects are orientation dependent^{12–16,35} and can often be interpreted in terms of a “rotational rainbow” effect, in which dependence of the potential energy surface on molecular orientation gives rise to one or more singularities in the classical distribution of angular momentum in the scattering product.

In diatomic molecule experiments, the observed rotational rainbow population distribution often resembles a Boltzmann distribution with an added Gaussian term, centered at the rainbow energy.^{18,36} In formaldehyde, there are three distinct projections of the angular momentum onto the molecular body frame and the problem is more complicated. Based on the results of our analysis in Section III B1, we attempt a model that

partitions the rotational energy into two parts: E_a (a -axis rotational energy) and E_{bc} (combined b - and c -axis rotational energy). We add a Gaussian contribution to the a -axis distribution function to obtain

$$P_i = Ng_i e^{-E_{bc,i}/kT_{bc}} (e^{-E_{a,i}/kT_a} + S e^{-(E_0 - E_{a,i})^2/\sigma_a^2}). \quad (1)$$

In eqn (1), P_i is the population of state i , N is a normalization factor, and g_i is a degeneracy factor that includes both M_J and nuclear spin state degeneracy. The first factor in eqn (1) describes the distribution of the E_{bc} partition of the energy, and the second factor describes the distribution of the E_a partition. T_a and T_{bc} are effective “temperatures” for the two types of energy and S parameterizes the extent to which population is distributed non-thermally in E_a . E_0 corresponds to the a -axis rotational energy of the rainbow, and σ_a is a width parameter that describes the sharpness of the rainbow. Note that eqn (1) converges to a Boltzmann distribution in the limit $T_a = T_{bc}$ and $S = 0$. Neglecting the relatively small contributions of asymmetry splitting and centrifugal distortion, we set

$$E_a = AK_a^2$$

$$E_{bc} = E_{\text{rot}} - AK_a^2,$$

where the rotational constant $A \approx 9.406$ cm^{-1} . In order to reduce correlation between the fit parameters, we constrain $T_a = T_{bc}$ (*i.e.* we used the same “Boltzmann temperature” for each degree of freedom), and we constrain the Gaussian width parameter to $\sigma_a = 850$ cm^{-1} , which is the value that gives the best fit at $E_i = 1.21$ eV, where the observed deviation from thermal distribution is greatest.

The spectrum was recorded several times at each E_i shown in Fig. 1, and each spectrum was fit individually to eqn (1). The average and standard deviation of the fit parameters obtained at each E_i are shown in Fig. 4(A–C) and are tabulated in Table SIII of the ESI.† Also shown in Fig. 4(D) is the percent improvement in



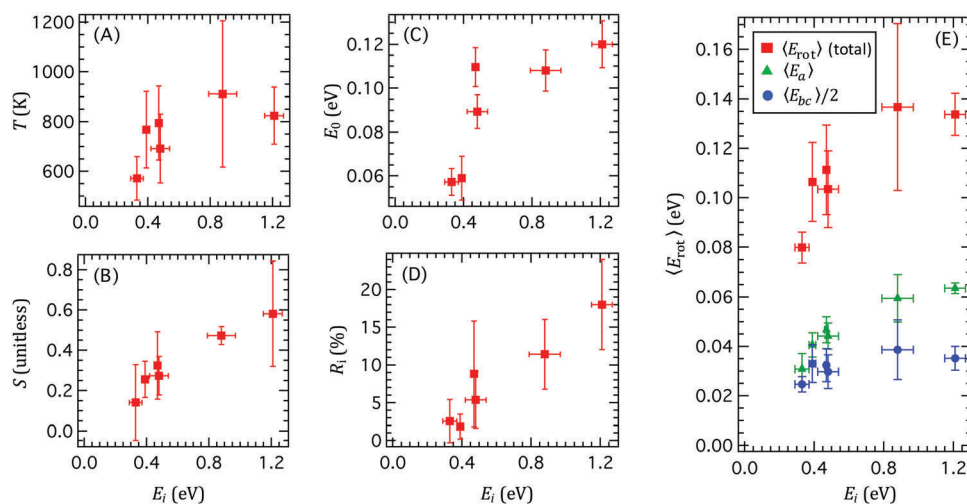


Fig. 4 The best-fit parameters obtained by fitting the spectra of scattered molecules to eqn (1), with the constraints $T_{bc} = T_a = T$ and $\sigma_a = 850 \text{ cm}^{-1}$, are shown as a function of incidence kinetic energy, E_i , in panels (A), (B) and (C). Panel (D) shows the average percent improvement, R_i , in the root mean square error of the fit, R_{fit} , relative to the RMSE obtained from a fit to a Boltzmann temperature, R_b . $R_i = (1 - R_b/R_{\text{fit}}) \times 100\%$. Panel (E) shows the expectation value, obtained from the fit model, for total rotational excitation, and for the components E_a and E_{bc} .

the root mean square error (RMSE) that is obtained from the fit to eqn (1), relative to the RMSE obtained from a fit to a Boltzmann distribution. Shown in Fig. 4(E) is the expectation value of rotational energy, $\langle E_{\text{rot}} \rangle$, obtained from the model at each incidence kinetic energy as well as the individual contributions, $\langle E_a \rangle$, and $\langle E_{bc} \rangle$. The best-fit spectrum obtained at each incidence energy is compared with the observed spectrum in Fig. 1, and the result obtained at $E_i = 1.21 \text{ eV}$ is included in Fig. 2 and 3.

Several trends are evident in the fit results. As E_i is increased, there is a slight trend towards higher T parameter (Fig. 4(A)), consistent with a general increase in both $\langle E_a \rangle$ and $\langle E_{bc} \rangle$. The average rotational energy obtained from the model for the scattered beam increases approximately linearly over the range $E_i = 0.33\text{--}0.8 \text{ eV}$ with a slope of $\langle E_{\text{rot}} \rangle \approx 0.24E_i$, and appears to exhibit some degree of saturation above $E_i = 0.8 \text{ eV}$ (Fig. 4(E)). In addition, the a -axis rainbow model reveals an increasing propensity for energy to be channeled into a -axis rotation as E_i is increased. This trend is evident from the increasing ratio between $\langle E_a \rangle$ and $\langle E_{bc} \rangle$ (Fig. 4(E)) and leads to a linearly increasing trend in the best-fit amplitude, S , of the non-thermal Gaussian term, which has a near-zero intercept at $E_i = 0$ (Fig. 4(B)). The fitted values for E_0 (Fig. 4(C)) indicate that the maximum in the non-thermal Gaussian contribution to the population distribution shifts from $K_a \approx 7$ at $E_i = 0.4 \text{ eV}$ to $K_a \approx 10$ at $E_i = 1.2 \text{ eV}$.

A consequence of the increasing importance of the non-thermal contribution to the population distribution is that the failure of the Boltzmann model to reproduce the observed spectrum increases strikingly as E_i is increased. At low incidence kinetic energy ($E_i < 0.4 \text{ eV}$), the observed spectrum is well reproduced by the Boltzmann fit, and the improvement of the fit error that is obtained by including the non-thermal Gaussian component of eqn (1) is negligible (Fig. 4(D)). However, at higher E_i , the Boltzmann fit fails to reproduce the spectral intensities (as shown in Fig. 2 and 3) and the improvement in the overall

RMSE obtained by including the Gaussian term reaches almost 20% at $E_i = 1.2 \text{ eV}$.

From a comparison of the observed spectra in Fig. 2 and 3 to the Boltzmann fit and the a -axis rainbow model fit, it is clear that the a -axis rainbow model provides significantly better agreement with experiment. The integrated intensities in the 1R_5 , 1R_7 , and 1R_9 bandheads obtained from the rainbow model all agree with experiment to within 16%, and the integrated intensity over the origins of the $^1R_{11-14}$ progressions ($28\,429\text{--}28\,435 \text{ cm}^{-1}$) agrees with experiment to within 4% (Fig. 2). The a -axis rainbow fit also correctly reproduces the observed relative intensities in the $^1P_{11}(11\text{--}16)$ lines between $28\,000$ and $28\,060$, and includes features from the 1P_9 branch such as the intensity due to $^1P_9(9)$ at $28\,110.6 \text{ cm}^{-1}$ (Fig. 3). Unlike the Boltzmann fit, the a -axis rainbow fit does not place too much intensity in the $^1P_1(33\text{--}38)$ lines or in the $^1P_{3,30}(33)$ line (discussed in Section III B1). Furthermore, the a -axis rainbow fit correctly reproduces (to within 0.05 cm^{-1}) the center frequencies of the observed features at $28\,076.0 \text{ cm}^{-1}$ and $28\,100.1 \text{ cm}^{-1}$, which are shifted relative to the Boltzmann fit due to contributions from $^1P_9(15)$ and $^1P_9(11)$.

In Fig. 5, the sum of the population in each K_a manifold (divided by the g_{ns} nuclear spin degeneracy) obtained from the fit to the a -axis rainbow model is plotted *versus* $E_a = AK_a^2$ as a function of E_i . At the lowest incidence energy, the population decreases approximately exponentially with E_a , as expected in a thermal distribution. As E_i is increased, a strong shoulder grows in at $\sim 0.12 \text{ eV}$, due to the increasing importance of the Gaussian term in eqn (1). The population in $K_a = 0$ is a factor of two below the trend because this manifold lacks the two-fold $k = \pm|K_a|$ degeneracy (*i.e.* there are no asymmetry doublets).

C. Classical scattering simulation

To aid in the interpretation of our results, we have calculated classical scattering trajectories. We treat the formaldehyde molecule as a rigid rotor with the geometry given by Burleigh *et al.*³⁷



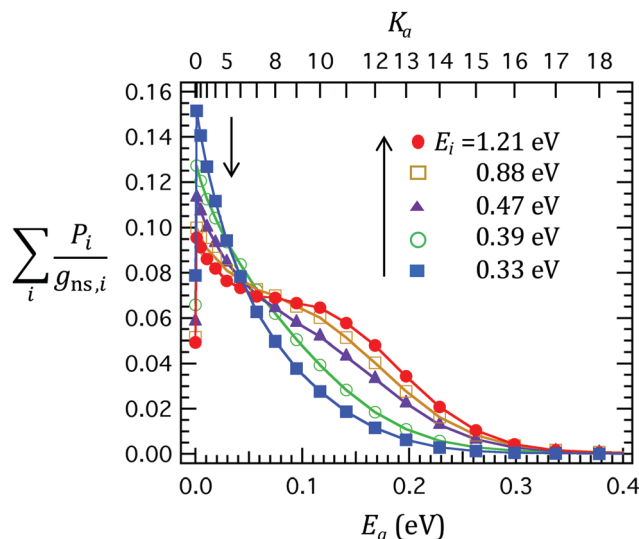


Fig. 5 The population within each K_a manifold, obtained from the a -axis rainbow model fit to the direct-scatter REMPI data is divided by the nuclear spin degeneracy (g_{ns}) and plotted as a function of E_i . The total population at each E_i is normalized to one. The population in $K_a = 0$ is a factor of two below the trend because this manifold lacks the two-fold $k = \pm|K_a|$ degeneracy.

We approximate the surface as a flat harmonic oscillator with mass 197 amu and the Debye frequency of gold ($\omega = k\Theta_D/\hbar$ with $\Theta_D = 170$ K). We tried two different simple models for the molecule–surface interaction. First, we used a purely repulsive hard-sphere potential (“hs”) with a hard wall at the van der Waals radius of each atom. Next, we used an empirical Lennard-Jones 6-12 potential (“L-J”), with parameters taken from the UFF force field.³⁸ The UFF van der Waals interaction underestimates the measured³⁰ and calculated^{39–41} binding energy of formaldehyde on Au(111) by more than an order of magnitude, so we scale the potential by a factor of 41 in order to bring the optimized binding energy into agreement with the measured value (0.31 eV).³⁰

The trajectories were calculated at normal incidence angle and random incidence orientation, over a range of incidence kinetic energies ($E_i = 0.33$ – 1.2 eV). The surface was initially at rest and the molecule was initially at a distance of $z_0 = 10$ Å from the surface with zero incident angular momentum. We integrated 10k trajectories for 2.4 ps at each E_i . For the L-J potential, any trajectory in which the molecule remained within 8 Å of the surface after 2.4 ps was labeled a “trapping” trajectory and was discarded. At $E_i = 0.33$ eV, 28% of trajectories were discarded on this basis. The proportion of “trapping” trajectories drops to 15% at $E_i = 0.47$ eV and to 2.1% at $E_i = 1.2$ eV. This trend moderately underestimates the observed trapping probability,³⁰ which drops from 57% at $E_i = 0.32$ eV to 25% at $E_i = 0.58$ eV.

After exiting the surface potential, the rotational motion of the directly scattered asymmetric top molecules propagates according to Newton’s equations of torque-free motion:

$$I_{aa}\dot{\omega}_a = \omega_b\omega_c(I_{bb} - I_{cc}) \quad (2a)$$

$$I_{bb}\dot{\omega}_b = \omega_c\omega_a(I_{cc} - I_{aa}) \quad (2b)$$

$$I_{cc}\dot{\omega}_c = \omega_a\omega_b(I_{aa} - I_{bb}). \quad (2c)$$

In the general case, eqn (2) leads to precession of the angular velocity about the three body-fixed internal axes, which “scrambles” the memory of the body-fixed components of the torque induced by the surface scattering event. However, for a near-prolate symmetric top, $I_{bb} - I_{cc} \approx 0$, and eqn (2a) reduces to $I_{aa}\dot{\omega}_a \approx 0$. As a consequence, ω_a (and $E_a = \frac{1}{2}\omega_a^2 I_{aa}$) are approximately conserved constants of motion, as is the quantity $E_{\text{rot}} - E_a = E_b + E_c$. However, the individual components E_b and E_c oscillate in time due to precession about the near-symmetry axis.

Fig. 6 shows the final average rotational energy about each internal axis of rotation, calculated from the two potentials as a function of E_i . The calculated values may be compared directly with the values from the experimental fits, shown in Fig. 4(E). Although the calculations overestimate the overall amount of rotational excitation, relative to the experimental fit, by a factor of ~ 2 , they nevertheless reproduce key qualitative features of the experimental observation. Most notably, regardless of whether an attractive potential is used, the calculation predicts that $\langle E_a \rangle$ excitation is greater than $\langle E_b \rangle$ or $\langle E_c \rangle$ excitation by an average factor of $2 \langle E_a \rangle / (\langle E_b \rangle + \langle E_c \rangle) = 1.6$ (L-J) or 1.3 (hs), compared to an experimental value of 1.5. When the purely repulsive hard sphere potential is used, rotational excitation increases linearly with E_i , but when the Lennard-Jones potential is used, the calculation also qualitatively reproduces the experimental observation that $\langle E_a \rangle$ exhibits some degree of saturation at higher E_i . These results suggest that the propensity to excite a -axis rotation is a simple consequence of the three-dimensional repulsive van der Waals shape of the formaldehyde molecule, whereas the saturation effect may be a signature of dynamical steering due to the orientation dependence of the attractive potential.

The top half of Fig. 7 shows the distribution of rotational energy about the internal axes of rotation calculated using the L-J potential with $E_i = 1.2$ eV. The E_b and E_c distributions can be approximated by an exponential decay. On the other hand, the E_a distribution exhibits a distinct non-exponential component, which gives rise to a broad feature between 0.15–0.4 eV. The calculated rotational distributions are in qualitative agreement

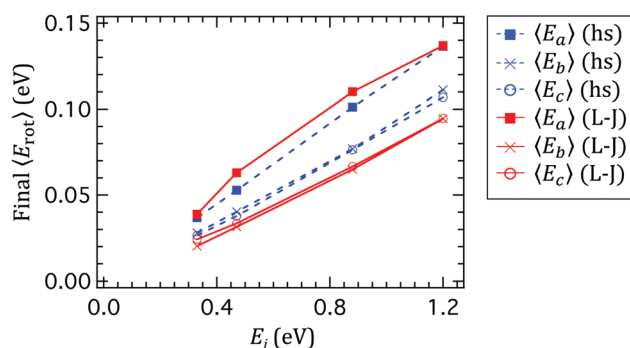


Fig. 6 The expectation value for rotational excitation about each principal internal axis of the formaldehyde molecule, obtained from classical scattering trajectory calculations, as a function of incidence kinetic energy, E_i , at normal scattering angle. Two interaction potentials were used: a repulsive hard-sphere potential (hs) and an empirically scaled Lennard-Jones potential (L-J). See text for further details.



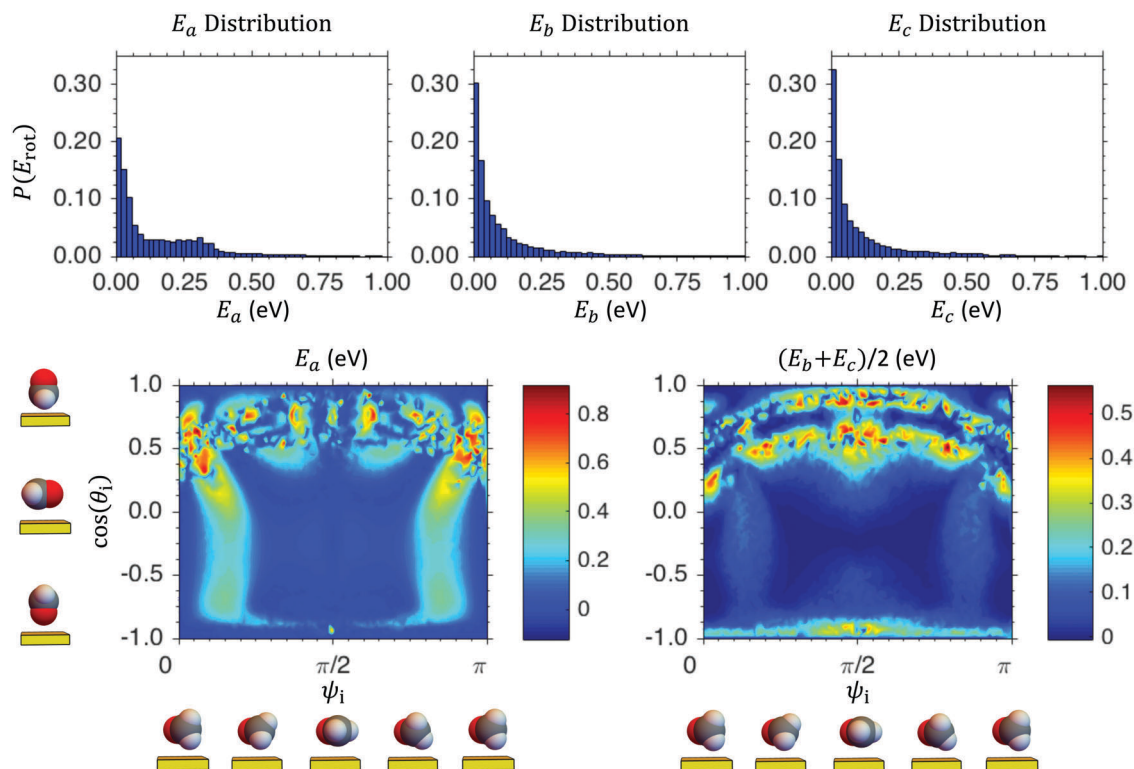


Fig. 7 Upper panels display the normalized distribution of rotational energy about each principal internal axis of the formaldehyde molecule, obtained from a classical scattering trajectory calculation using the L-J potential at $E_i = 1.2$ eV, at normal scattering angle. The bottom panels show the rotational energy components, E_a and $(E_b + E_c)/2$, as a function of incidence orientation, described by the Euler angles θ_i and ψ_i . See text for further details.

with the distributions obtained from our fit model. Namely, the E_b and E_c distributions resemble an exponential Boltzmann population distribution whereas the E_a distribution resembles an exponential decay plus an added Gaussian component (see Fig. 5).

The classical trajectory calculation is not in complete quantitative agreement with the experimental fit model. The calculation overestimates the average overall rotational excitation, and the calculated non-Boltzmann hump in the E_a distribution is broader and extends to higher energy than that of the fit model. The trajectory calculation places the center of the hump at 0.30 eV whereas the fit model places the center at 0.12 eV. However, since the trajectory calculation uses only a crude empirical potential energy surface, it is unlikely to capture the orientation dependence of the potential accurately and complete quantitative agreement cannot be expected. Theoretical investigations involving a more sophisticated *ab initio* potential energy surface are merited, but are beyond the scope of the current work. Nevertheless, since the classical trajectories reproduce key qualitative features of the experiment, they provide clues as to the likely origin of these features.

The bottom half of Fig. 7 shows the components, E_a and $(E_b + E_c)/2$, of rotational energy, calculated as a function of the incident Euler angles, θ_i and ψ_i . In our chosen coordinate system, θ_i represents the polar angle between the molecular CO-bond symmetry axis and the surface normal, and ψ_i represents the angle between the surface normal and its orthogonal

projection onto the CH_2 plane. See cartoons in Fig. 7 for clarification. Normal angle scattering of a C_{2v} molecule from a flat, structureless surface is totally symmetric in the molecule-fixed axis frame with respect to the third Euler angle, ϕ , which is omitted from the figure. The b - and c -axis projections of the rotational energy are maximally excited at incidence orientations where the CO axis is nearly (but not exactly) along the surface normal ($\theta_i \approx 0$ or π), and are minimally excited when the incident CO axis orientation is “flat-on” ($\theta_i \approx \pi/2$). At incidence angles near $\theta_i \approx 0$ (CH_2 group oriented toward the surface), the prevalence of “multi-bounce” collisions, where the two hydrogen atoms strike the surface in rapid succession, give rise to complicated structure in all three components of E_{rot} . Apart from a relatively narrow region near $\theta_i = \pi$, the plot of $(E_b + E_c)/2$ versus incident orientation does not reveal any widespread regions over which the function exhibits a smooth maximum.

In contrast, E_a exhibits broad regions of incidence orientation space over which it is excited to approximately the same value. Most strikingly, when $\pi/20 < \psi_i < \pi/4$ or $3\pi/4 < \psi_i < 19\pi/20$, excitation to $E_a \approx 0.3$ eV is highly likely, regardless of the value of θ_i . The explanation is simple. A large degree of a -axis excitation results whenever one of the hydrogens strikes the surface at a skew angle of ψ_i , regardless of the polar angle θ_i between the CO axis and the surface normal. This occurs even at incident orientations in which the CH_2 group is initially pointed away from the surface ($\theta_i < 0$) due to “double-bounce” collisions in which the O and CH_2 moieties strike the surface in rapid succession.



Since this behavior arises over a wide swath of orientation space, excitation probability accumulates at $E_a \approx 0.3$ eV over many trajectories and gives rise to the distinctly non-Boltzmann propensity for high K_a excitation which is observed. This explanation allows us to interpret the effect in terms of a sterically induced rotational rainbow scattering phenomenon. Because there is a wide region of orientation space over which the Jacobian $\partial E_a / \partial(\psi_i, \theta_i)$ is approximately zero (and a long curve over which $\partial E_a / \partial \psi_i$ is rigorously zero), the classical probability distribution for E_a as a function of ψ_i contains singularities for excitation to the local extrema in E_a .

IV. Discussion

A. Possible sources of error in the analysis

Alternative explanations for the apparent non-Boltzmann intensities must be considered and excluded. In ref. 32 we characterized our REMPI scheme and found no significant discrepancies between our measured relative intensities and the relative intensities in the calculated absorption spectrum, to within an uncertainty of $\sim 20\%$. Although the spectra analyzed in that work were obtained at rotational temperatures of 300 K or less, and did not include significant intensity from transitions involving states with $K_a'' \geq 9$, our results allow us to rule out the possibility that the observed intensity at higher K_a'' arises from artifacts such as K_a -dependent ionization cross sections or K_a -dependent dissociative lifetimes of the intermediate \tilde{A} state levels. The intensities from $K_a'' = 9$ transitions measured in the trapping/desorption channel (top spectrum, Fig. SI, ESI†) agree with the Boltzmann simulation to within 10%. Furthermore, the magnitude of the observed non-thermal relative intensities in the direct scatter channel grows in as a function of E_i , and is therefore consistent with a population effect and not with incorrect relative line intensities. In addition, the high- K_a'' populations that one obtains through analysis of the ${}^{\text{R}}K_a''$ branches matches that which one obtains from the ${}^{\text{P}}K_a''$ branches, although these transitions involve \tilde{A} -state intermediate levels with different K_a' . The results that we report by fitting the entire band are reproduced when either the ${}^{\text{R}}$ or ${}^{\text{P}}$ region is fitted independently.

Another possible source of intensity error in our measurement arises from the fact that we have not performed exhaustive rotationally-resolved measurement of the speed distribution of scattered molecules. Therefore, our population estimates are based on scattered beam density measurements that have not been converted to flux, and are only valid in the limit $\Delta v / \langle v \rangle \ll 1$ (*i.e.* when the velocity spread is small compared with the mean velocity.) Experimental limitations make it difficult to perform precise rotationally resolved time-of-flight measurements. However, we have made coarse speed measurements for a representative sample of rotational states differing in E_{rot} and K_a . The results, which are detailed in Section SV of the ESI,† provide an upper limit for $\Delta v / \langle v \rangle < 10\%$ for the rotational state dependence of the average scattered velocity at $E_i = 1.21$ eV, which is far too small to account for deviation of the observed intensities from a Boltzmann model.

We have also investigated the effect of laser polarization, and we find no difference in relative intensities when the dye laser polarization is rotated from horizontal to vertical polarization. Orientation of the angular momentum due to the surface is likely scrambled due to nuclear hyperfine dephasing before the scattered molecules reach the detection region.

B. Trapping probability at high E_i

Shown in the top panel of Fig. 8 is the measured trapping probability of formaldehyde on the Au(111) surface as a function of incidence energy at normal incidence angle, reported in ref. 30. Following Fig. 6 of ref. 30, the trapping probabilities from three other weakly bound, non-activated scattering systems (with binding energies in the range 0.18–0.33 eV) of similar molecular mass (28–30 amu) and surface atom mass (195–197 amu) are shown for comparison. In a hard cube model,^{42–44} these systems are expected to exhibit similar trapping probability. However, in comparison with the diatomic molecules NO and CO, formaldehyde exhibits a significantly higher trapping probability. The shift of ~ 0.3 eV in the initially decaying portion of the curve is much too large to be explained by differences in binding energy. Strikingly, formaldehyde exhibits 25% trapping probability at an incidence energy of 0.58 eV—more than an order of magnitude higher than the trapping probability of NO at similar incidence energies. This result indicates that formaldehyde is able to dissipate much larger amounts of incident kinetic energy during collision with the Au(111) surface. The polyatomic prolate top molecule ethane exhibits a similar degree of enhanced trapping in the range $E_i = 0.05$ –0.4 eV.

In classical trajectory simulations of ethane trapping on Pt(111) reported in ref. 4, the excitation of molecular rotation was found to play a decisive role in the trapping probability, especially for the highest normal incidence kinetic energy investigated (0.4 eV). In trajectories where the first bounce led to sufficient excitation of molecular rotation, the molecule became temporarily “rotationally trapped” in the surface potential. The average energy transferred to phonons during the first bounce was the similar in the “rotational trapping” and “single-bounce scattering” channels. In $>90\%$ of “rotationally trapped” trajectories, the initially stored rotational energy was dissipated to surface degrees of freedom during subsequent bounces, leading eventually to a relaxed trapping configuration. Thus, the degree of rotational excitation that occurred during the first bounce determined the eventual fate of the molecule (trapping or direct scattering) to within 10% uncertainty. In ref. 30, it was remarked that a similar mechanism might explain the trapping of formaldehyde on Au(111) at $E_i = 0.58$ eV.

We examine the feasibility of the “rotational trapping” mechanism by analysis of our experimentally measured rotational distribution. Assuming that the average phonon excitation during the first bounce is determined by a hard sphere collision model involving a single gold atom, a 30 amu formaldehyde molecule will lose 46% of its incident energy to phonon excitation.³⁰ At $E_i = 0.58$ eV, an additional 0.31 eV of kinetic energy must therefore, on average, be dissipated in order for trapping to occur. If we make the (admittedly naive)



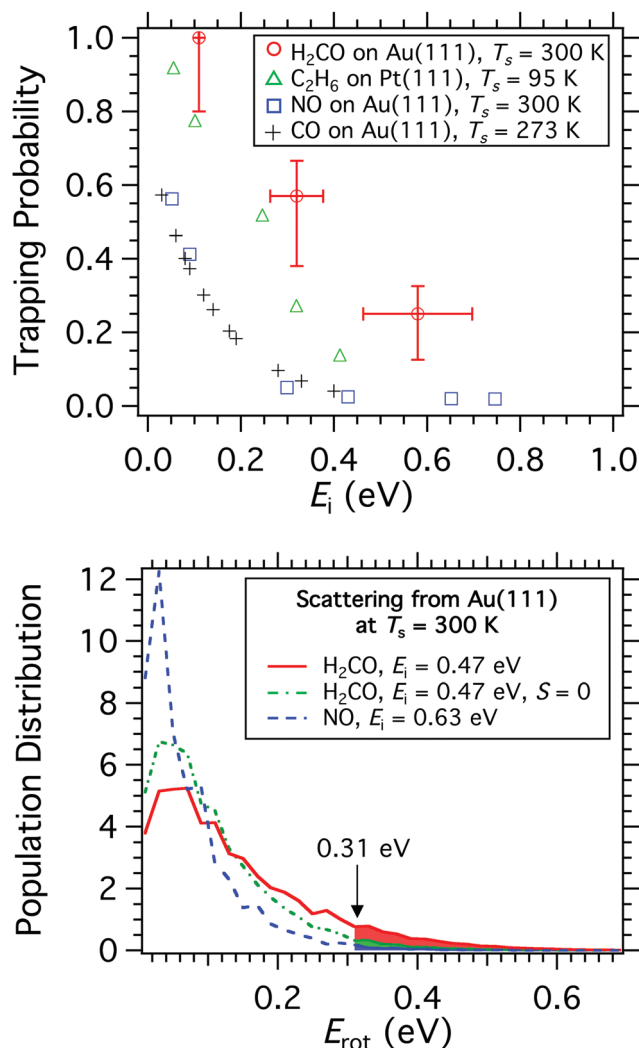


Fig. 8 In the top panel, the estimated trapping probability of formaldehyde on $\text{Au}(111)$ ³⁰ is plotted as a function of incidence kinetic energy. Measured trapping probabilities from three other weakly bound molecular scattering systems with similar molecular and surface atom masses are shown for comparison. Data for ethane on $\text{Pt}(111)$ (0.33 eV binding energy) is taken from ref. 46; NO on $\text{Au}(111)$ (0.24 eV binding energy) is from ref. 47; and CO on $\text{Au}(111)$ (0.18 eV binding energy⁴⁸) is from ref. 49. The bottom panel shows the normalized rotational energy distribution that results from direct scatter of formaldehyde at $E_i = 0.47$ eV. Shown for comparison are the distribution that is obtained when the S parameter, describing the contribution of the a -axis rotational rainbow, is set to zero. Also shown is the distribution that is obtained from the direct scatter of NO from $\text{Au}(111)$ at $E_i = 0.63$ eV, reported in ref. 45. The tails of the distribution with $E_{\text{rot}} \geq 0.31$ eV are shaded. This is the portion of the distribution with sufficient rotational excitation to give rise to rotationally mediated trapping, according to a simple model (see text).

assumption that there is no correlation between energy transfer to phonons and to molecular rotation, then the trapping probability that will result from rotational excitation mechanisms will be given approximately by the fraction of molecules with ≥ 0.31 eV of rotational excitation. In our fitted rotational distribution, shown in the bottom panel of Fig. 8, 9.8% of directly scattered formaldehyde molecules at $E_i = 0.47$ eV are excited to this energy. This is less than the observed trapping

probability of 25% (at $E_i = 0.58$ eV), but is obtained at a slightly lower value of E_i and represents only the rotational energy distribution in the direct scatter channel (*i.e.*, in molecules that *do not* trap). The semi-quantitative agreement is therefore a strong argument that rotation plays a major role in the trapping mechanism. On the other hand, in the ground state rotational distribution of NO molecules scattered from $\text{Au}(111)$ at $E_i = 0.60$ eV (shown for comparison in Fig. 8) only 1.1% of molecules are excited to $E_{\text{rot}} \geq 0.31$ eV.⁴⁵ This is consistent with the observation that the trapping probability in the NO on $\text{Au}(111)$ system is an order of magnitude less than that of formaldehyde. Finally, we note that without the a -axis rainbow (*i.e.*, with the S parameter of eqn (1) set to zero) the fraction of directly scattered formaldehyde molecules with ≥ 0.31 eV of rotational excitation is only 3.1% at $E_i = 0.47$ eV. (This distribution is also shown for comparison in Fig. 8) This underscores the importance of a -axis rotation to the high-energy tail of the rotational distribution and points to the likely contribution of this additional degree of rotational freedom (not present in diatomic molecules) to trapping at higher incidence kinetic energies.

V. Conclusions

In this work, we have used a $1 + 1'$ REMPI scheme to characterize the ground state rotational distribution that results from molecular beam surface scattering of formaldehyde from $\text{Au}(111)$ in both the trapping/desorption and direct scatter regime. Our work is the first demonstration of rotational distribution measurements from surface scattering of an asymmetric top molecule. In the direct scatter channel, there is a pronounced propensity to excite a high degree of a -axis (twirling) rotation, consistent with a high- J_a axis-specific rotational rainbow. Classical trajectory simulations allow the result to be interpreted as a steric effect, arising from excitation to high J_a over a wide range of incident orientation space over which $\partial J_a / \partial \psi_i = 0$. We note that a -axis rotation makes a significant contribution to the high-energy tail of the rotational distribution and may therefore play an important role in the trapping mechanism at high incidence energy. Since the rotational rainbow arises primarily from steric effects and inertial considerations, one can expect such effects to occur quite generally in the scattering of weakly bound, small prolate (or near-prolate) top molecules in which the rotation of light, off-axis hydrogen atoms is readily excited.

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References

- 1 C. T. Rettner, C. B. Mullins, D. S. Bethune, D. J. Auerbach, E. K. Schweizer and W. H. Weinberg, Molecular beam studies of trapping dynamics, *J. Vac. Sci. Technol., A*, 1990, **8**(3), 2699–2704.
- 2 M. A. Hines and R. N. Zare, The interaction of CO with Ni(111): rainbows and rotational trapping, *J. Chem. Phys.*, 1993, **98**(11), 9134–9147.
- 3 D. Kulginov, M. Persson, C. Åkerlund, I. Zorić and B. Kasemo, CO₂ sticking on Pt(111): the role of kinetic energy and internal degrees of freedom, *J. Vac. Sci. Technol., A*, 1995, **13**(3), 1511–1516.
- 4 J. A. Stinnett, R. J. Madix and J. C. Tully, Stochastic simulations of the trapping of ethane on Pt(111) from a realistic potential: the roles of energy transfer processes and surface corrugation, *J. Chem. Phys.*, 1996, **104**(8), 3134–3142.
- 5 J. F. Weaver and R. J. Madix, Trapping dynamics of isobutane, *n*-butane, and neopentane on Pt(111): effects of molecular weight and structure, *J. Chem. Phys.*, 1999, **110**(21), 10585–10598.
- 6 R. Schlögl, Heterogeneous catalysis, *Angew. Chem., Int. Ed.*, 2015, **54**(11), 3465–3520.
- 7 F. Zaera, Probing catalytic reactions at surfaces, *Prog. Surf. Sci.*, 2001, **69**(1–3), 1–98.
- 8 R. R. Cavanagh and D. S. King, Rotational- and spin-state distributions: NO thermally desorbed from Ru(001), *Phys. Rev. Lett.*, 1981, **47**, 1829–1832.
- 9 M. C. Lin and G. Ertl, Laser probing of molecules desorbing and scattering from solid surfaces, *Annu. Rev. Phys. Chem.*, 1986, **37**, 587–615.
- 10 J. Kimman, C. T. Rettner, D. J. Auerbach, J. A. Barker and J. C. Tully, Correlation between kinetic-energy transfer to rotation and to phonons in gas-surface collisions of NO with Ag(111), *Phys. Rev. Lett.*, 1986, **57**, 2053–2056.
- 11 L. V. Novakoski and G. M. McClelland, Orientation of CHF₃ desorbed and scattered from Ag(111): measurements using electrostatic focusing, *Phys. Rev. Lett.*, 1987, **59**, 1259–1262.
- 12 L. Vattuone, L. Savio, F. Pirani, D. Cappelletti, M. Okada and M. Rocca, Interaction of rotationally aligned and of oriented molecules in gas phase and at surfaces, *Prog. Surf. Sci.*, 2010, **85**(1–4), 92–160.
- 13 M. G. Tenner, F. H. Geuzebroek, E. W. Kuipers, A. E. Wiskerke, A. W. Kleyn, S. Stolte and A. Namiki, Orientation dependence of rotational excitation in NO scattering from Ag(111), *Chem. Phys. Lett.*, 1990, **168**(1), 45–50.
- 14 F. H. Geuzebroek, A. E. Wiskerke, M. G. Tenner, A. W. Kleyn, S. Stolte and A. Namiki, Rotational excitation of oriented molecules as a probe of molecule-surface interaction, *J. Phys. Chem.*, 1991, **95**(21), 8409–8421.
- 15 A. W. Kleyn, Rotational rainbow scattering of oriented molecules, *Surf. Rev. Lett.*, 1994, **1**(1), 157–173.
- 16 N. Bartels, K. Golibrzuch, C. Bartels, L. Chen, D. J. Auerbach, A. M. Wodtke and T. Schäfer, Dynamical steering in an electron transfer surface reaction: oriented NO($v = 3$, $0.08 < E_i < 0.89$ eV) relaxation in collisions with a Au(111) surface, *J. Chem. Phys.*, 2014, **140**(5), 054710.
- 17 A. W. Kleyn and T. C. M. Horn, Rainbow scattering from solid surfaces, *Phys. Rep.*, 1991, **199**(4), 191–230.
- 18 A. W. Kleyn, A. C. Luntz and D. J. Auerbach, Rotational energy transfer in direct inelastic surface scattering: NO on Ag(111), *Phys. Rev. Lett.*, 1981, **47**, 1169–1172.
- 19 R. Schinke, Rotational rainbows in diatom(solid) surface scattering, *J. Chem. Phys.*, 1982, **76**(5), 2352–2359.
- 20 H. Voges and R. Schinke, On the dynamics of molecule/surface scattering, *Chem. Phys. Lett.*, 1983, **95**(3), 221–225.
- 21 C. W. Muhlhausen, L. R. Williams and John C. Tully, Dynamics of gas-surface interactions: scattering and desorption of NO from Ag(111) and Pt(111), *J. Chem. Phys.*, 1985, **83**(5), 2594–2606.
- 22 J. A. Barker, A. W. Kleyn and D. J. Auerbach, Rotational energy distributions in molecule surface scattering: model calculations for NO/Ag(111), *Chem. Phys. Lett.*, 1983, **97**(1), 9–13.
- 23 J. C. Polanyi and R. J. Wolf, Dynamics of simple gas-surface interaction. II. Rotationally inelastic collisions at rigid and moving surfaces, *J. Chem. Phys.*, 1985, **82**(3), 1555–1566.
- 24 D. C. Jacobs and R. N. Zare, Simplified trajectory method for modeling gas-surface scattering: the NO/Pt(111) system, *J. Chem. Phys.*, 1989, **91**(5), 3196–3207.
- 25 M. E. Coltrin and B. D. Kay, Quasiclassical trajectory study of rotational energy transfer in the scattering of NH₃ from a flat, rigid gold surface, *J. Chem. Phys.*, 1988, **89**(1), 551–561.
- 26 B. G. Perkins, Jr. and D. J. Nesbitt, Quantum-state-resolved CO₂ scattering dynamics at the gas-liquid interface: incident collision energy and liquid dependence, *J. Phys. Chem. B*, 2006, **110**(34), 17126–17137.
- 27 K. Golibrzuch, J. H. Baraban, P. R. Schirhatti, J. Werdecker, C. Bartels and A. M. Wodtke, Observation of translation-to-vibration excitation in acetylene scattering from Au(111): a REMPI based approach, *Z. Phys. Chem.*, 2015, **229**(10–12), 1929–1949.
- 28 A. C. Wight and R. E. Miller, Rainbow scattering of methane from LiF(100): probing the corrugation and anisotropy of the gas-surface potential, *J. Chem. Phys.*, 1998, **109**(5), 1976–1982.
- 29 B. D. Kay, T. D. Raymond and M. E. Coltrin, Observation of a dynamical propensity rule in rotationally inelastic gas-surface scattering: NH₃ on Au(111), *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1987, **36**, 6695–6697.
- 30 B. C. Krüger, G. Barratt Park, S. Meyer, R. J. V. Wagner, A. M. Wodtke and T. Schäfer, Trapping-desorption and direct-scattering of formaldehyde at Au(111), *Phys. Chem. Chem. Phys.*, 2017, DOI: 10.1039/c7cp03907g.
- 31 G. Barratt Park, B. C. Krüger, S. Meyer, D. Schwarzer and T. Schäfer, The ν_6 fundamental frequency of the \tilde{A} state of formaldehyde and Coriolis perturbations in the $3\nu_4$ level, *J. Chem. Phys.*, 2016, **144**(194308).
- 32 G. Barratt Park, B. C. Krüger, S. Meyer, A. M. Wodtke and T. Schäfer, A $1 + 1'$ resonance-enhanced multiphoton ionization scheme for rotationally state-selective detection of formaldehyde via the $\tilde{A}^1A_2 \leftarrow \tilde{X}^1A^1$ transition, *Phys. Chem. Chem. Phys.*, 2016, **18**, 22355–22363.



- 33 D. J. Clouthier and D. A. Ramsay, The spectroscopy of formaldehyde and thioformaldehyde, *Annu. Rev. Phys. Chem.*, 1983, **34**, 31–58.
- 34 H. S. P. Müller, G. Winnewisser, J. Demaison, A. Perrin and A. Valentin, The ground state spectroscopic constants of formaldehyde, *J. Mol. Spectrosc.*, 2000, **200**(1), 143–144.
- 35 E. W. Kuipers, M. G. Tenner, A. W. Kleyn and S. Stolte, Observation of steric effects in gas-surface scattering, *Nature*, 1988, **334**(6181), 420–422.
- 36 J. C. Tully and M. J. Cardillo, Dynamics of molecular motion at single-crystal surfaces, *Science*, 1984, **223**(4635), 445–450.
- 37 D. C. Burleigh, A. B. McCoy and E. L. Sibert, An accurate quartic force field for formaldehyde, *J. Chem. Phys.*, 1996, **104**(2), 480–487.
- 38 A. K. Rappe, C. J. Casewit, K. S. Colwell, W. A. Goddard III and W. M. Skiff, UFF, a full periodic table force field for molecular mechanics and molecular dynamics simulations, *J. Am. Chem. Soc.*, 1992, **114**(25), 10024–10035.
- 39 W.-K. Chen, S.-H. Liu, M.-J. Cao, Q.-G. Yan and C.-H. Lu, Adsorption and dissociation of methanol on Au(111) surface: a first-principles periodic density functional study, *J. Mol. Struct.: THEOCHEM*, 2006, **770**(1–3), 87–91.
- 40 S. Liu, P. Jin, D. Zhang, C. Hao and X. Yang, Reaction mechanism for methanol oxidation on Au(111): a density functional theory study, *Appl. Surf. Sci.*, 2013, **265**, 443–451.
- 41 L. Wang, C. He, W. Zhang, Z. Li and J. Yang, Methanol-selective oxidation pathways on Au surfaces: a first-principles study, *J. Phys. Chem. C*, 2014, **118**(31), 17511–17520.
- 42 R. M. Logan and J. C. Keck, Classical theory for the interaction of gas atoms with solid surfaces, *J. Chem. Phys.*, 1968, **49**(2), 860–876.
- 43 E. W. Kuipers, M. G. Tenner, M. E. M. Spruit and A. W. Kleyn, Differential trapping probabilities and desorption of physisorbed molecules: application to NO/Ag(111), *Surf. Sci.*, 1988, **205**(1–2), 241–268.
- 44 C. R. Arumainayagam, M. C. McMaster, G. R. Schoofs and R. J. Madix, Dynamics of molecular CH₄ adsorption on Pt(111), *Surf. Sci.*, 1989, **222**(1), 213–246.
- 45 R. Cooper, Z. Li, K. Golibrzuch, C. Bartels, I. Rahinov, D. J. Auerbach and A. M. Wodtke, On the determination of absolute vibrational excitation probabilities in molecule-surface scattering: case study of NO on Au(111), *J. Chem. Phys.*, 2012, **137**(6), 064705.
- 46 C. R. Arumainayagam, G. R. Schoofs, M. C. McMaster and R. J. Madix, Dynamics of molecular adsorption of ethane with platinum(111): a supersonic molecular beam study, *J. Phys. Chem.*, 1991, **95**(3), 1041–1047.
- 47 A. M. Wodtke, H. Yuhui and D. J. Auerbach, Insensitivity of trapping at surfaces to molecular vibration, *Chem. Phys. Lett.*, 2005, **413**(4–6), 326–330.
- 48 D. P. Engelhart, R. J. V. Wagner, A. Meling, A. M. Wodtke and T. Schäfer, Temperature programmed desorption of weakly bound adsorbates on Au(111), *Surf. Sci.*, 2016, **650**, 11–16.
- 49 C. T. Rettner, The search for direct vibrational excitation in gas-surface collisions of CO with Au(111), *J. Chem. Phys.*, 1993, **99**(7), 5481–5489.

