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State-to-state methane-surface scattering as a probe of catalytic activity

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Quantum state-resolved scattering experiments for methane molecules colliding with a catalytically active nickel surface are compared to scattering from a nickel surface passivated by a single layer of graphene. The vibrational state distribution of the scattered methane is observed to differ dramatically for the two surfaces. Quantum-mechanical inelastic scattering calculations show that these differences are related to the catalytic activity of the surface impact site. Our results demonstrate how inelastic scattering can be used to probe the reactive potential-energy surfaces of molecule-metal systems important to heterogeneous catalysis.

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I. INTRODUCTION

Uncovering the dynamics of a chemical reaction between a gas-phase molecule and a solid surface is key to understanding and predicting heterogeneous catalysis, a process of great economic and ecological value. For example, solid-phase catalysts are involved in the production of the fuel that powers the combustion engine in our automobiles as well as in cleanup of its exhaust gases. The search for more efficient catalysts is therefore of great economic and ecological importance and can undoubtedly benefit from a detailed, microscopic understanding of the dynamical processes occurring on catalytically active surfaces.

Most studies of the reaction dynamics at the gas-surface interface have focused on reactive events by detecting reaction products and measuring how reaction rates depend on the energy and internal state of incident molecule [1,2]. Consider the dissociation of methane on transition-metal catalysts; of particular interest since it is the rate-limiting step in steam reforming, a process of great importance for the chemical industry. During the last two decades, this reaction has been extensively explored, both experimentally and computationally, revealing nonstatistical dynamics that include mode specificity [3], bond selectivity [4,5], and steric effects [6]. However, much less is known about the fate of those molecules that fail to react during their collision with the catalyst surface and instead scatter back inelastically into the gas phase. This is especially true for polyatomic molecules such as methane because of a lack of generally applicable and highly sensitive detection techniques that can provide information on the vibrational and rotation quantum states of

the scattered species. The lack of scattering data is unfortunate because it leaves us blind to important features of the interactions governing reactions. As we demonstrate in this paper, those state distributions carry signatures of the rich dynamics that unfold as the incident molecule makes its way along the reaction path, before being repelled back into the gas phase. By contrast, molecules that succeed in reacting on the surface lose all memory of those dynamics as they interact with the plethora of surface states.

Recently, we have implemented a state-resolved method for detecting molecules scattered from surfaces based on a cryogenic bolometer and quantum state-specific infrared laser tagging [7,8]. The method, which we call bolometer infrared laser tagging (BILT), enables the state-resolved detection of methane or any molecule with an infrared-active vibrational mode and a rotationally resolved infrared spectrum. The combination of quantum state-specific preparation of the reactant molecules in the incident molecular beam by infrared laser pumping, with state-resolved BILT detection of the scattered species, enables so-called “state-to-state” scattering experiments. They provide highly detailed information about the scattering dynamics and energy transfer between the molecular and surface degrees of freedom. When state-to-state scattering experiments are accompanied by a quantum-dynamics simulation on a realistic potential energy hypersurface calculated through density-functional theory, it becomes possible to develop a complete and detailed picture of the different inelastic and reactive processes that control the outcome of a molecule/surface collision.

Here, we report dramatic changes in the product state distribution observed for state-to-state scattering of methane from clean and graphene-covered Ni(111) surfaces. The experimental findings are compared to fully quantum-mechanical scattering calculations, which relate the amount of intramolecular vibrational energy redistribution (IVR) to the catalytic activity of the impact site; i.e., the ability of a particular surface site to activate, or break, a C–H bond. We thus demonstrate how inelastic scattering can be used to probe the reactive potential-energy

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surfaces of molecule–metal systems important to heterogeneous catalysis.

II. METHODS

The state-to-state scattering experiments were carried out in a previously described [9,10] molecular beam–surface science apparatus. Briefly, a continuous molecular beam of methane was produced by supersonic expansion of CH₄ seeded in He. Quantum state-specific preparation of the surface incident methane was achieved by infrared laser excitation of the CH₄ molecules in the molecular beam using a tunable continuous-wave optical parametric oscillator system (IR-OPO). The state-prepared methane collided with a clean or graphene-covered Ni(111) sample in an ultrahigh-vacuum chamber. Scattered methane molecules were detected with quantum state resolution by a highly sensitive liquid He-cooled bolometer in combination with state-specific laser tagging using a second IR-OPO system by virtue of the BILT technique, as we have demonstrated in a recent publication [8]. Details of the experimental methods are described in the Supplemental Material [11].

To examine the vibrationally inelastic scattering of CH₄ we use a quantum approach that has successfully described the dissociative chemisorption of methane on Ni and Pt surfaces [12–15]. This method, which provides a very accurate treatment of the molecular vibrations, is described in these earlier works and the Supplemental Material [11], and we summarize only the essential ideas here.

We consider normal incidence and treat the center-of-mass motion of CH₄ parallel to the surface (coordinates X and Y) using a sudden approximation. That is, we compute the scattering probabilities for impact on a particular surface site, keeping X and Y fixed, then repeat this for several other impact sites, averaging the results. This has worked well for similar systems [16,17]. For each impact site we locate the minimum energy path (MEP) as the molecule moves towards the surface. To compute the energy we use VASP, the Vienna *Ab initio* Simulation Package, developed at the Institut für Materialphysik of the Universität Wien [18–22]. Exchange-correlation effects are included using a Specific Reaction Parameter functional shown to accurately describe the dissociative chemisorption of methane on Ni and Pt surfaces [23].

At several dozen points along each MEP we compute the total energy $V_0(s)$, where s is the distance along the path in mass-weighted coordinates. We also compute and diagonalize the Hessian to find the normal vibrational coordinates Q_k and corresponding frequencies, $\omega_k(s)$, describing motion orthogonal to the MEP in the harmonic approximation. Ignoring anharmonic terms we write our potential-energy surface, for a given MEP, in the reaction path coordinates s and Q_k , as [24]

$$V = V_0(s) + \sum_{k=1}^9 \frac{1}{2} \omega_k^2(s) Q_k^2. \quad (1)$$

This harmonic treatment is accurate for the strongly bound stretching and bending vibrations of the methane, modes 1 through 9, but it is not a good description of X , Y , or rotational motion. As noted, X and Y are treated with a sudden model. The rotational degrees of freedom are allowed to evolve adi-

abatically, reasonable at these low energies. The remaining 10 degrees of freedom, s and the Q_k , are described by expanding our molecular wave function in a vibrational basis [11–13,15,24]:

$$\Psi(t) = \sum_{\mathbf{n}} \chi_{\mathbf{n}}(s;t) \Phi_{\mathbf{n}}(\{Q_k\}; s). \quad (2)$$

The $\Phi_{\mathbf{n}}$ are vibrationally adiabatic harmonic oscillator eigenfunctions that depend parametrically on s , and \mathbf{n} labels each vibrational state. As we time evolve the wave packets, $\chi_{\mathbf{n}}$, nonadiabatic couplings create transitions between different vibrationally adiabatic states. The backscattered wave packets are analyzed to extract energy–resolved vibrational state probabilities [12].

III. RESULTS

Figure 1 shows a comparison of two BILT spectra recorded when state-prepared CH₄($\nu_3 = 1$, $J = 1$) is scattered from a bare Ni(111) and from a graphene-covered Ni(111) surface, respectively. The quantum-state population of the scattered CH₄ is encoded in these spectra with each line corresponding to a vibrational transition originating from a rovibrationally excited quantum state that was populated by the molecule–surface collision. In order to record BILT spectra such as the ones shown in Fig. 1, the infrared tagging laser was scanned over a spectral region covering both $\nu_3 \rightarrow 2\nu_3$ and $\nu_1 \rightarrow \nu_1 + \nu_3$ hot-band transitions [11]. Vibrationally excited states at lower energy than ν_1 were also probed and found to contain no detectable population irrespective of which of the two surfaces was present.

There is striking difference between the BILT spectra obtained for collisions with bare Ni(111) and those with Gr/Ni(111). When scattering from the bare Ni(111) surface [8], we detect both vibrationally elastic scattering into the ν_3 vibrational state as well as vibrationally inelastic scattering into the ν_1 vibrational state, which is approximately 100 cm^{−1} lower in vibrational energy than ν_3 . On the other hand, when scattering from Gr/Ni(111), all lines coming from the ν_1 vibrational state are absent and we detect only vibrationally elastic scattering into different rotational levels of the ν_3 vibrational state. This indicates that a single layer of graphene deposited on the Ni(111) surface completely eliminates the $\nu_3 \rightarrow \nu_1$ vibrational energy redistribution that was easily detected with a branching ratio of 40% for scattering from the bare Ni(111) surface [8].

Our quantum dynamics simulations described below confirm that the absence of surface-induced IVR is due to changes in the CH₄–metal interaction when a single layer of graphene is deposited on top of the Ni(111) surface. Furthermore, the simulations suggest that the propensity for $\nu_3 \rightarrow \nu_1$ IVR scales with the chemical reactivity of the surface site where the scattering takes place.

Besides information on surface-induced IVR, the recorded spectra also contain information on the molecule–surface rotation energy transfer during the direct scattering event. The incident CH₄($\nu_3 = 1$, $J = 1$) is rotationless in the laboratory frame since its total angular momentum ($J = 1$) is solely due to vibrational angular momentum ($l = 1$) intrinsic in the triply degenerate ν_3 vibration. Following scattering from the

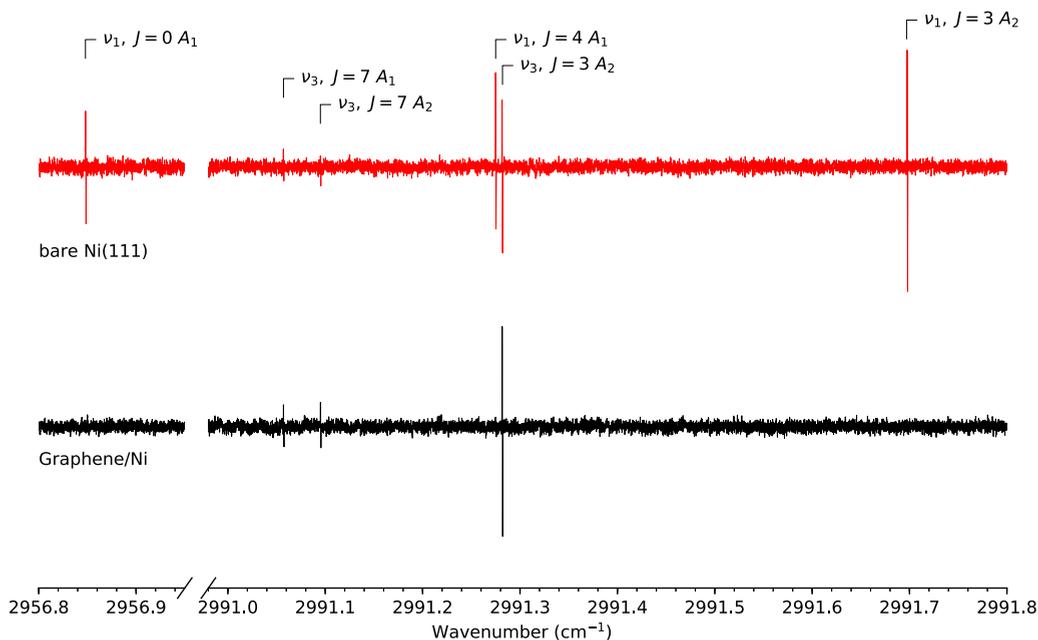


FIG. 1. Examples of BILT spectra obtained for scattering of state-prepared $\text{CH}_4(\nu_3; J = 1)$ from clean Ni(111) (top trace in red) and graphene-covered Ni(111) (lower trace in black). Each spectral line marks a specific rovibrational quantum state of the scattered CH_4 as indicated by the labels. For scattering from Ni(111), 40% of the incident $\text{CH}_4(\nu_3; J = 1)$ molecules undergo $\nu_3 \rightarrow \nu_1$ surface-induced vibrational redistribution (IVR). For scattering from Gr/Ni(111), $\nu_3 \rightarrow \nu_1$ surface-induced IVR is absent with all the scattered CH_4 remaining in the ν_3 state indicated by the fact that all lines that originate from the ν_1 state are missing from the lower spectrum. (The spectral lines have a derivative form due to the use of frequency modulation for lock-in detection of the bolometer signal.)

bare Ni(111) surface, state-resolved BILT detection was used to quantify the rotational-state population with J up to 8 as shown in Fig. 2.

Figures 2 and 3 show how for the same incident kinetic energy, scattering from Gr/Ni(111) leads to stronger rotational excitation than scattering from bare Ni(111). Such a difference in rotational excitation could be caused by the larger corrugation of Gr/Ni(111) compared to Ni(111) but

other explanations such as differences in physisorption well depths cannot be excluded without theoretical modeling. Figure 3 shows rotational Boltzmann plots for the rotational distributions of Fig. 2. Except for the lowest J levels, which correspond to (near)-elastic scattering, the rotational-state populations can be fitted by a Boltzmann distribution with a rotational temperature of 153 ± 8 K for Ni(111) and 208 ± 14 K for Gr/Ni(111), much lower than the 673 K temperature

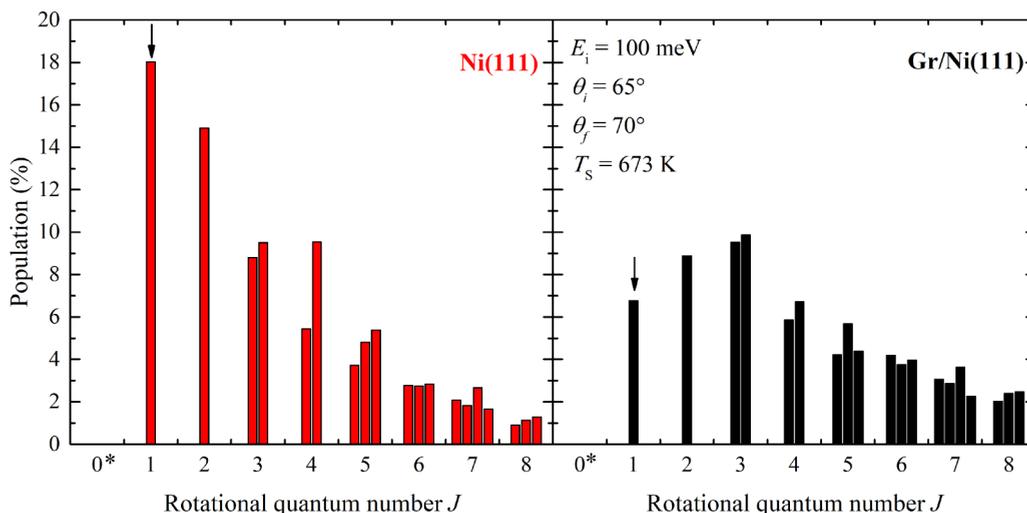


FIG. 2. Rotational level populations for $\text{CH}_4(\nu_3, J)$ scattered from Ni(111) and Gr/Ni(111). The incident CH_4 was prepared in the $(\nu_3 = 1, J = 1)$, state which is labeled by an arrow \downarrow . For $J > 2$, multiple bars give the population of the individual sublevels for each value of J . Stronger rotational excitation is observed for scattering from Gr/Ni(111) compared to bare Ni(111). (* For symmetry reasons, the $J = 0$ level does not exist for the nuclear spin conformer of methane prepared in the incident beam.)

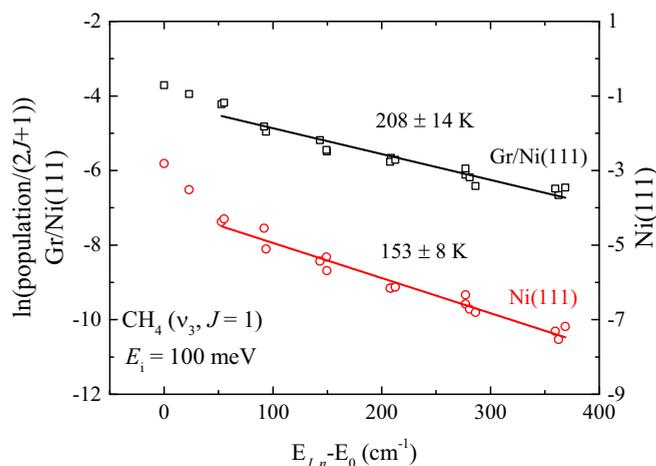


FIG. 3. Boltzmann fits for the rotational level populations shown in Fig. 2 as a function of rotational energy $E_{J,n}$ of the scattered CH_4 for incident kinetic energy $E_i = 100$ meV and a surface temperature of 673 K. E_0 is the energy of the lowest rotational state $J = 1$. Note the use of different vertical scales for Gr/Ni(111) (left) and Ni(111) (right) for clarity of presentation. Solid lines are linear fits to the data excluding the two lowest rotational states $J = 1$ and 2. The characteristic rotational temperatures given by the slopes of the linear fits are 153 ± 8 K for Ni(111) and 208 ± 14 K for Gr/Ni(111).

of the target surface. The fact that both rotational temperatures are far below the surface temperature and that the scattering leads to nearly specular angular distributions [11] confirms that we are probing a fast and direct scattering process rather than a trapping-desorption channel in which case the incident molecules would thermalize with the surface before desorbing back into the gas phase.

Our calculations consider collisions at 16 points on the surface unit cell, reducible to 4 unique impact sites: the top, bridge, half-bridge, and near hollow sites [11]. Results for each site are shown in Fig. 4 for the case of a rigid lattice. We find that the $\nu_3 \rightarrow \nu_1$ transition probability is the largest by far, that there is a strong variation in vibrational inelasticity with impact site, and that this correlates with the catalytic activity of the impact site. Density-functional theory (DFT) studies of methane dissociation on Ni and Pt surfaces find that the highest catalytic activity is over the top site [25], and that the barrier increases as we move towards the bridge site [15]. While we are well below these barriers at the incident energies considered here, our DFT calculations show that as the CH_4 approaches a reactive site, the C–H bond closest to the surface starts to elongate. The lower the barrier at a particular site, the sooner this increase in C–H bond length starts to happen with respect to the distance above the surface.

In Fig. 4 we list the length of this C–H bond for each impact site, at the point where the carbon is at a distance Z_C above the surface such that $V_0(Z_C) = 0.2$ eV; i.e., at the point of closest approach for a collision energy of 0.2 eV. We clearly see that the amount of C–H stretching correlates with both catalytic activity and the magnitude of the $\nu_3 \rightarrow \nu_1$ transition. This makes sense, since we are distorting the molecule close to the turning point, and this distortion, roughly a C–H stretch, mixes the stretch normal-mode vibrations ν_3 and ν_1 , which

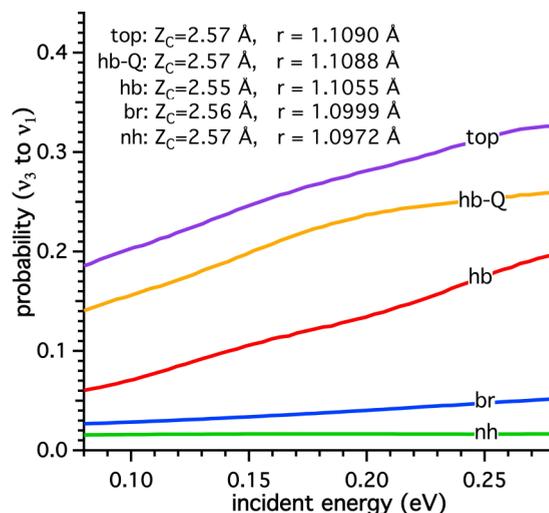


FIG. 4. Computed probabilities for the $\nu_3 \rightarrow \nu_1$ transition as a function of energy, for impact at the top, bridge (br), half-bridge (hb), and near-hollow (nh) sites, and at the half-bridge site where the Ni atom is puckered by 0.1 \AA (hb-Q). Also shown are the distance of the carbon above the surface when $V_0 = 0.2$ eV, Z_C , and the length of the reactive C–H bond at this distance, r .

(classically) differ only by the relative phases of the C–H stretching motion of the four bonds. At these low collision energies, we do not observe any bending distortions of the molecule near the turning point. Thus, we see only very small transition probabilities to the $2\nu_2$, $2\nu_4$ and $\nu_2 + \nu_4$ bending states. A recent calculation of H_2O scattering from Cu(111) found a similar mixing of the two stretching vibrations [26]. While our study relates IVR in polyatomic-surface scattering with catalytic activity, a paper published over 30 years ago [27] predicted that the lengthening of the bond during a low-energy collision of H_2 with a catalytically active surface could induce rovibrational transitions. Since then, a few other studies have noted how the existence of a dissociation channel can modify nonreactive diatom-surface scattering [28–30]. Finally, note that the site-to-site variation in the classical turning point, Z_C , is very small at 0.2 eV, and that the largest value of Z_C is for the catalytically active top site. Thus the observed effects are not related to how close the molecule gets to the metal.

Lattice displacement changes the barriers to methane dissociation on metals, leading to strong temperature effects in dissociative chemisorption [31,32]. In Fig. 4 we plot results for impact at the half-bridge site, where the Ni atom closest to the methane is puckered out of the plane of the surface by 0.1 \AA , roughly the rms vibrational amplitude of a Ni atom at 300 K. On Ni(111), this puckering lowers the barrier by about 0.1 eV [25], and we see that this lattice distortion significantly increases the $\nu_3 \rightarrow \nu_1$ transition probability and the C–H bond length, just as it increases the catalytic activity.

To more rigorously include these lattice motion effects, we would need to repeat these calculations, at all sites, for several values of the surface atom displacements, and then Boltzmann- and site average the results. However, we can estimate the total $\nu_3 \rightarrow \nu_1$ transition probability for a rigid surface by averaging over the impact sites with their proper

weighting [11], and we find that it is about 10% at 0.2 eV. As in the experiments, the variation with energy is relatively weak for this near-resonant nonactivated transition, particularly given acceleration by the 0.2 eV attractive well. By including lattice motion at the high temperatures of the experiment, our results in Fig. 4 suggest that this could easily increase by a factor of 2 or 3, giving a branching ratio very similar to that in the experiments. A more quantitative comparison with experiment would require a better treatment of rotation, and perhaps also X and Y , since translational steering may be important at these low collision energies. Such a full 15-dimensional quantum treatment is not currently feasible.

To test our assertions, we repeat our calculations for the case where the Ni(111) surface is covered with a graphene overlayer. The methods for modeling this in a supercell DFT calculation are described elsewhere [33]. We considered methane impact at two sites: directly over a C atom that is directly over a Ni atom, and over the graphene hollow, which is over the Ni(111) hexagonal close-packed hollow. The graphene layer is kept rigid. The methane-Gr/Ni(111) well depth is 0.26 eV at both sites. We find that the C–H bonds of methane are not elongated at all, suggesting that the surface is catalytically inactive, as expected. Moreover, at a collision energy of 0.15 eV, the $\nu_3 \rightarrow \nu_1$ transition probability at both of these sites is only about 1.5%, compared to 25% for the (bare) Ni(111) top site. These findings are consistent with our scattering experiments.

In conclusion, our state-to-state CH₄ scattering experiments from bare and graphene-covered Ni(111) show that surface-induced IVR from the antisymmetric to the symmetric C–H stretch normal mode occurs on the bare Ni(111) but is completely absent if a single layer of graphene is added to passivate the Ni(111) surface. Quantum scattering calculations reveal that the propensity for the observed $\nu_3 \rightarrow \nu_1$ vibrational energy redistribution process is related to the extent of C–H bond stretching during the molecule's closest approach to a scattering site, which in turn correlates with the catalytic activity of the specific surface site. Combined, the results of our experimental and theoretical work indicate that the probing of surface-induced IVR processes by state-to-state scattering experiments of polyatomic molecules from metal surfaces yields valuable information on the reactive potential-energy surface and the catalytic activity of the molecule–surface system under study. Extensions of this work to a wider range of experimental conditions are currently underway.

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