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Vibrational energy relaxation of interfacial OH on a water-covered α -Al₂O₃(0001) surface: a non-equilibrium *ab initio* molecular dynamics study

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Vibrational relaxation of adsorbates is a sensitive tool to probe energy transfer at gas/solid and liquid/solid interfaces. The most direct way to study relaxation dynamics uses time-resolved spectroscopy. Here we report on a non-equilibrium *ab initio* molecular dynamics (NE-AIMD) methodology to model vibrational relaxation of OH vibrations on a hydroxylated, water-covered α -Al₂O₃(0001) surface. In our NE-AIMD approach, after exciting selected O–H bonds their coupling to surface phonons and to the water adlayer is analyzed in detail, by following both the energy flow in time, as well as the time-evolution of Vibrational Density of States (VDOS) curves. The latter are obtained from Time-dependent Correlation Functions (TCFs) and serve as prototypical, generic representatives of time-resolved vibrational spectra. As most important results, (i) we find a few-picosecond lifetime of the excited modes and (ii) identify both hydrogen-bonded aluminols and water molecules in the adsorbed water layer as main dissipative channels, while the direct coupling to Al₂O₃ surface phonons is of minor importance on the timescales of interest. Our NE-AIMD/TCF methodology is powerful for complex adsorbate systems, in principle even reacting ones, and opens a way towards time-resolved vibrational spectroscopy.

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1. Introduction

Water-covered metal oxides play an important role in many areas of surface science, with applications ranging from heterogeneous catalysis over electrochemistry to geochemistry.¹ Tremendous effort has been investigated in recent years to characterize various water/metal oxide interfaces,^{2–9} which improved our molecular-level understanding of their surface physical chemistry. Vibrational spectroscopy is a powerful tool to do so. Notably, surface-sensitive Vibrational Sum Frequency (VSF) spectroscopy has been employed for silica/water,¹⁰ quartz/water,¹¹ or alumina (Al₂O₃)/water¹² interfaces. Here we concentrate on the latter, as a particularly well studied model system for insulating metal oxides.

While most of the vibrational studies above are of the steady-state type, a more direct account of dynamics is provided by time-resolved spectroscopy, *e.g.*, VSF in a pump–probe mode.¹³ Time-resolved spectroscopy can give direct information on energy transfer between an adsorbate and the surface, which is central for the

understanding of adsorbate bonding but also for more applied issues, such as energy storage or bond-selective chemistry. In the pump–probe variant of VSF spectroscopy,¹⁴ a spectrally narrow IR (infrared) pulse is used to excite/pump an adsorbate vibration, which is then probed by two spatially and temporarily overlapping IR and visible pulses which generate a VSF signal that is recorded as a function of frequency and the delay time Δt , between pump and probe pulses. Both steady-state and time-resolved VSF spectra have recently been recorded for water-covered α -Al₂O₃(0001) surfaces by Borguet *et al.*,^{15–17} for example, focusing primarily on the hydrogen-bonded spectral region and on the effects of interfacial electrolytes on O–H stretching vibrations.

In our recent work, in a combined experimental/theoretical effort, the so-called Al–I terminated α -Al₂O₃(0001) surface^{18–22} as the most stable alumina surface in vacuum,^{19–22} further a hydroxylated α -Al₂O₃(0001),^{21,23} as well as O-terminated α -Al₂O₃(1 $\bar{1}$ 02)²⁴ and O-terminated α -Al₂O₃(11 $\bar{2}$ 0)²⁵ surfaces, all in contact with small amounts of water were studied with steady-state VSF spectroscopy, mostly to characterize the vibrations of interfacial O–H (or O–D) bonds. From the theory side, periodic Density Functional Theory (DFT) was used, either stationary and together with Normal Mode Analysis (NMA),^{20,21,24,25} or in the form of *ab initio* molecular dynamics (AIMD) in combination with Time-dependent Correlation Function (TCF) methods,^{22,23} to characterize adsorbate vibrations. The AIMD/TCF

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approach is computationally costly but also very powerful, as it accounts for Vibrational Density Of States (VDOS) curves and even for spectra, IR and VSF,^{22,23} beyond the harmonic approximation and it explicitly includes the effects of temperature, which allows for direct comparison between simulations and experiments.

The motivation for our present, theoretical work is to extend the previous AIMD/TCF simulations explicitly to the time-dependent domain, by initially vibrationally exciting specific O–H chromophores of a water-covered alumina surface, and follow their dynamics after increasing time delays, in the spirit of a pump–probe vibrational analysis. Specifically, we shall consider a water-covered, hydroxylated α -alumina interface (see below), and excite high-frequency O–H vibrations associated with non-hydrogen bonded aluminol groups, *i.e.*, quasi-free O–H units attached to surface Al ions. In fact, this surface was extensively studied by us before,^{21,23} and currently corresponding pump–probe VSF experiments are underway to study vibrational relaxation.²⁶ We shall try, on the basis of non-equilibrium *ab initio* molecular dynamics (NE-AIMD) simulations, to (i) model the vibrational energy relaxation times of excited surface aluminols in $\text{H}_2\text{O}/\alpha\text{-Al}_2\text{O}_3(0001)$ and (ii) understand the vibrational relaxation mechanism and energy flow in this system. (iii) A third goal for this work is to extend the known NE-AIMD methodology,²⁷ to calculate time-resolved vibrational spectra for complex interfaces. For that, NE-AIMD will be combined with time-windowed TCF methods, with time-dependent correlation functions serving to compute spectra $I(\omega, \Delta t)$ as needed for interpreting pump–probe IR/VSF signals, for example. Here, we shall not make the full step to pump–probe IR/VSF spectra, though, and compute time-resolved VDOS curves instead, *i.e.*, time-resolved power spectra that can be seen as a generic precursors to real spectra. On top, the VDOS curves help to analyze the relaxation process, complementary to other tools, such as kinetic energy analysis.

Concerning the vibrational energy redistribution it is common practice to define a “vibrational lifetime”, as the population decay time of the excited chromophore’s $\nu = 1$ state to the ground state, $\nu = 0$, *i.e.*, $P_{\nu=1}(t) \propto \exp(-t/\tau)$. Note that this definition holds strictly for quantum mechanics and for $T = 0$ only and in fact, neither in experiment nor in our classical theory as here one necessarily operates with $P_{\nu=1}(t)$ curves but uses related measures instead. For excited O–H stretching vibrations at non-metallic interfaces, a wide range of lifetimes can be expected. In case when the OH bond was part of a hydrogen-bonded network, *e.g.*, a water layer, time-resolved VSF experiments for the air/water interface gave lifetimes ≤ 1 ps.^{14,28} Here the excited interfacial OH relaxes efficiently thanks to its direct coupling with H_2O bending overtones, which then allows further relaxation into the hydrogen bonding modes. If, on the other hand, an isolated O–H stretch mode, say, was directly coupled to surface phonons only, one could expect much longer lifetimes. In this case the high-frequency O–H mode can only relax *via* slow multi-phonon processes, because the Debye frequency of the surface is much lower. For α -alumina, the Debye frequency is in the order of 1000 cm^{-1} , so three to four phonons would be required for relaxation at least. Multi-phonon relaxation is indeed a slow

process: an extreme example is the C–O stretch mode of carbon monoxide on $\text{NaCl}(100)$, with $\tau \sim 4 \text{ ms}$.²⁹ Another example, chemically closer to α -alumina is represented by the O–H stretching relaxation of interfacial silanols in silica. For this system, early time-resolved vibrational spectroscopy measurements^{30,31} indicated a lifetime in the order of 100 ps. In the present study, the excited O–H chromophores are neither strongly H-bonded nor in contact with surface phonons only (there are water molecules around, see below), so an intermediate situation may be expected. In fact, we shall find lifetimes in the order of several picoseconds below.

Our paper is organized as follows. In the next Section II, the model of the interface studied here, hydroxylated $\alpha\text{-Al}_2\text{O}_3(0001)$ with additional water will be introduced and static DFT and canonical AIMD calculations on this system done in ref. 23 will be reviewed that are of relevance also for the present study. In Section III, the NE-AIMD methodology used here will be introduced, analysis methods will be described together with the NE-AIMD/TCF approach to time- and frequency resolved VDOS spectra. In Section IV we present results, *i.e.*, (i) lifetime calculations based on kinetic-energy analysis, (ii) subsystem-resolved energy flows to unravel the relaxation mechanism, (iii) time- and frequency resolved VDOS spectra for lifetime estimates, and as the precursors for time-resolved vibrational spectra just mentioned. A final Section V summarizes and concludes this work.

II. The surface model, static DFT and canonical AIMD calculations

A. The water/aluminol interface model

As the pump–probe VSF experiments²⁶ were done on a $\alpha\text{-Al}_2\text{O}_3(0001)$ surface sample exposed to air or to liquid water at room temperature, we employed a surface model that roughly represents α -alumina at low/medium water coverage. The model is the same as used in ref. 23 for the interpretation of vibrational spectra of water-covered α -alumina(0001). It consists of a hydroxylated alumina surface which forms from the so-called Al-terminated, UHV-stable (0001) surface.^{18,32–34} The latter has an atomic-layer sequence Al–O₃–Al–Al–O₃–Al–R, where “R” stands for a repetition of the Al–O₃–Al sequence into the bulk (when referring to a 1×1 cell). When exposed to water, the top layer Al-atoms are removed and the topmost O layer is saturated with hydrogen – the hydroxylated surface (HS) has formed. The atomic-layer sequence of the HS is (HO)₃–Al–(Al–O₃–Al)–R. Within a 2×2 cell, which we use here, we have twelve terminating OH bonds, or aluminols. We consider in addition, five Al layers and two oxygen layers, giving a simulation cell of composition $\text{Al}_{20}\text{O}_{36}\text{H}_{12}$. In our model, on top of the upper surface layer we adsorb eight intact water molecules. We refer to this system as the “Hydroxylated Surface +2 MonoLayers” (2ML) model, or HS + 2ML as in ref. 23. The HS + 2ML structure was found particularly stable by Density Functional Theory (DFT) calculations, due to the formation of a honeycomb-like layer network consisting of water hexagons.²³

The high stability is due to the formation of hydrogen bonds between some of the aluminol OH groups and molecular H_2O .

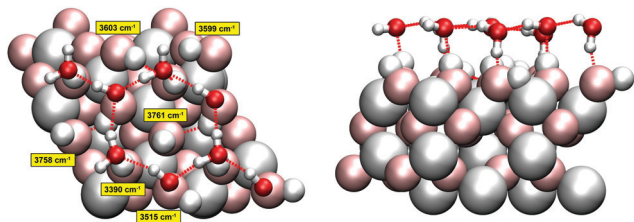


Fig. 1 Top (left) and side views (right) of the hydroxylated α - $\text{Al}_2\text{O}_3(0001)$ surface (2×2), with additional eight water molecules (HS + 2ML). Our model has a stoichiometric composition of 92 atoms ($\text{Al}_{20}\text{O}_{36}\text{H}_{12} \cdot (\text{H}_2\text{O})_8$). Surface atoms are shown as “van der Waals spheres” in pale colors, water thin films in a “ball-and-stick” representation (H in white, Al in grey and O in red, surface atoms large, water atoms small). Hydrogen bonds are indicated by dashed red lines. In the left panel, for some of the O–H groups harmonic frequencies, calculated at the PBE + D2 level of theory, are indicated in wavenumbers. In particular, there are four non-hydrogen-bonded aluminol OH groups, two of them “upright” and with very high frequencies (3758 cm^{-1} and 3761 cm^{-1}), and two of them in “parallel” orientation with lower frequencies (3603 cm^{-1} and 3599 cm^{-1}). Even lower frequencies are found for hydrogen-bonded aluminols – two examples are shown. In AIMD, only the uppermost three atomic layers (an O layer and a double-layer of Al) plus the adsorbate atoms were allowed to move, while the lower layers were kept frozen at PBE + D2 bulk geometries.²¹

In the model of Fig. 1, four of the twelve aluminol OH groups are non-hydrogen bonded (“quasi-free”), while eight are involved in H-bonds. The water layer appears to be also very rigid during AIMD trajectories, which makes this distinction between hb- (hydrogen-bonded) and nhb-aluminols (non-hydrogen bonded) valid on all timescales considered here and in ref. 23. As seen in Fig. 1, the quasi-free aluminols can come in “upright” and more “parallel” orientations relative to the surface. We note that in geometry optimization, NMA and AIMD only the uppermost three atomic layers plus the hydrogens of our HS + 2ML were allowed to move, while the lower layers were kept frozen at bulk geometries.^{21,23} The same protocol was followed in the AIMD simulations for the present work.

B. Stationary DFT and canonical AIMD calculations

For the HS + 2ML model shown in Fig. 1, we performed geometry optimization, NMA, and AIMD simulations,²³ all at the Kohn–Sham³⁵ DFT (KS-DFT) level of theory, employing the Perdew–Burke–Ernzerhof (PBE)³⁶ exchange–correlation functional and a projector-augmented plane-wave basis (PAW)³⁷ as implemented in the Vienna Ab initio Simulation Package (VASP).^{38–40} Dispersive van der Waals interactions were included by Grimme’s empirical D2 method.⁴¹ For all calculations a $3 \times 3 \times 1$ k -point mesh and a kinetic energy cutoff of 400 eV were used. The 2×2 cell slab models used have an extension of $9.66 \text{ \AA} \times 9.66 \text{ \AA} \times 31.41 \text{ \AA}$ (20 \AA of vacuum in the z -direction) and employ 3D periodic boundary conditions.

In ref. 23, besides NMA for harmonic frequency analysis, we also modelled vibrational spectra (Vibrational Densities of States, VDOS, IR and VSF) of the hydroxylated α -alumina(0001) surface without (HS) and with (HS + 2ML) additional water, using AIMD at 300 K with the Nosé–Hoover thermostat.^{42–44} Specifically, we fully employed surface-specific Velocity–Velocity Autocorrelation Functions (VVAFs) to calculate ssp-polarized VSF

spectra.⁴⁵ AIMD showed that the aluminol OH bonds on the hydroxylated (0001) surface, are orientationally flexible at room temperature: within a ps-timescale, they can switch from the “upright” position (polar angle (θ_{OH}) less than 45°) to a “parallel” one, where the bonds are almost flat with respect to the (0001) surface plane. With an additional water layer, this flexibility is somewhat hindered. Further details about the orientational behaviour of aluminols OH can be found in ref. 23. In all AIMD calculations a timestep of 0.2 fs was adopted.

In ref. 23, the following characteristic frequencies/frequency ranges for HS + 2ML were found by NMA:

- parallel aluminols show vibrational frequencies around 3600 cm^{-1} if not involved in accepting hydrogen bonds from nearby water molecules;
- non-hydrogen bonded upright aluminols have higher vibrational frequencies, in the range of $3750\text{--}3760 \text{ cm}^{-1}$;
- hydrogen-bonded aluminol OH and water–OH groups are both red-shifted with respect to the quasi-free OH groups, to a range between $3000\text{--}3500 \text{ cm}^{-1}$;
- water bending modes are found around $1600\text{--}1660 \text{ cm}^{-1}$;
- below $\sim 1100 \text{ cm}^{-1}$, one finds H_2O librational modes, aluminol OH bending, substrate phonons and water layer vibrations, respectively.

For illustration, we show in Fig. 1, left, some selected aluminol vibrational frequencies. Special attention will be given to non-hydrogen bonded aluminol OH groups, two of them “upright” and with very high frequencies (3758 cm^{-1} and 3761 cm^{-1}) in the static model, and two of them in “parallel” orientation with lower frequencies (3603 cm^{-1} and 3599 cm^{-1}). From the AIMD vibrational spectra at room temperature, which contain anharmonicity and temperature effects, rather similar values are obtained – mainly broadened w.r.t. to NMA stick spectra.²³ In particular the high-frequency modes are only slightly red-shifted relative to the NMA frequencies, by a few wavenumbers: the non-hydrogen bonded aluminol OH vibrations give rise to a VDOS peak centered at slightly under 3750 cm^{-1} , with a Full Width at Half Maximum (FWHM) of around 35 cm^{-1} (see ref. 23 and below). Our theoretical observations are in agreement with published experimental VSF spectra.^{12,21} (It should also be noted, however, that other works see only weak or no signals in this high-frequency range for water-covered α -alumina(0001) surfaces and α -alumina particles.⁴⁶) Within our model, at an IR excitation frequency of around 3700 cm^{-1} , we can reasonably expect that mainly the quasi-free aluminol OH groups are excited. Further, due to selection rules for IR light polarized perpendicular to the surface, notably the “upright” non-hydrogen bonded OH bonds should be excitable.

III. Non-equilibrium NE-AIMD and NE-AIMD/TCF approaches

A. AIMD calculations after vibrational excitation by “velocity swapping”

In order to model the IR preexcitation and subsequent relaxation, measured by time-resolved VSF, we employ a combination of equilibrium and non-equilibrium AIMD methods. Specifically, a

protocol illustrated in Fig. 2 is followed, consisting of the following steps:

(1) Starting geometry and equilibration phase: a geometry optimization at the PBE + D2 level of theory is done for the initial HS + 2ML structure. From there, five canonical *NVT*/AIMD trajectories at 300 K are run, each 24 ps long.

(2) Post-equilibration phase: starting from the end points of phase (1), five microcanonical *NVE*/AIMD trajectories are run, first for 10 ps, then for another 10 ps. The post-equilibration phase is introduced to remove spurious effects caused by coupling with the thermostat. The two different end times of phase (2) are used to test the influence of the length of the post-equilibration phase. We will loosely refer to phase (2) also as the “equilibrium *NVE*” phase.

(3) IR excitation by “velocity swapping”, subsequent non-equilibrium dynamics: the two end times of phase (2) (10 ps and 20 ps after equilibration phase (1)), are used as starting times for two sets of non-equilibrium AIMD simulations after “IR excitation” (*NEQ*₁, *NEQ*₂), called also “non-equilibrium *NVE*” in what follows. Both for *NEQ*₁ and *NEQ*₂, five *NVE*/AIMD trajectories are run after “swapping” of nuclear velocities of specific O–H bonds, all 10 ps long.

(4) Analysis: data are analyzed for energy relaxation.

Let us specify stage (3) of the scheme in Fig. 2, where atom velocities of selected O–H bonds associated with interfacial aluminols are “swapped”.⁴⁷ This step is used to mimic an experimental IR excitation in the pump–probe scheme. The interaction of a single vibrational chromophore with IR light involves the absorption of a vibrational quantum of energy, $\hbar\omega_0$. In our AIMD scheme, this excess energy is assumed to be fully converted into kinetic energy of atoms involved in excited aluminols. In this way, swapping of atom velocities enables us to perform non-equilibrium simulations, without the need of more complex techniques (*e.g.*, explicit coupling to an IR pulse). The “kinetic energy-only model” used here does not include

any atom displacements initially, along (unknown) vibrational coordinates. Of course, the excess energy stored in the O–H bonds at the beginning of phase (3), will rapidly and periodically be transferred into potential energy, and then be redistributed into other modes. For further features of the original equilibrium/non-equilibrium AIMD model, see ref. 27, 47 and 48.

The velocity swapping itself proceeds as follows. First, as an idealization we only swap velocities of H- and O-atoms associated with an excited aluminol. This is achieved by choosing, as swapped velocities $\tilde{v}(0)$ at time $t = 0$ (the time at the end of phase (2)):⁴⁷

$$\tilde{v}_H(0) = v_H(0) + \varepsilon_H \hat{L}_{OH}(0) \quad (1)$$

$$\tilde{v}_O(0) = v_O(0) + \varepsilon_O \hat{L}_{OH}(0) \quad (2)$$

Here,

$$\varepsilon_H = \frac{(p - p_0)}{m_H}; \quad \varepsilon_O = -\frac{(p - p_0)}{m_O}$$

$$p_0 = \mu_{OH}(v_H(0) - v_O(0)) \cdot \hat{L}_{OH}(0); \quad p = (p_0^2 + 2\mu_{OH}\Delta E)^{1/2}.$$

In the equations above, $\mu_{OH} = \frac{m_H m_O}{m_H + m_O}$ is the O–H bond reduced mass and $\hat{L}_{OH}(0) = \frac{r_{OH}(0)}{|r_{OH}(0)|}$ the unit vector along the OH bond. The excess energy is represented by ΔE and in our simulations this value is taken as $\hbar\omega_0 = 3750 \text{ cm}^{-1}$ per excited bond, in order to account for our computed frequencies (see above). Velocities $v(0)$ are those prior to swapping.

Only selected aluminol OH bonds are pre-excited in our AIMD simulations. Specifically, due to IR frequency and dipole orientation issues as described above, we excite only non-hydrogen bonded, “upright” aluminols. This is achieved in practice by analyzing at the end of phase (2), polar angles θ_{OH} of all non-hydrogen-bonded aluminols. Then, if $\theta_{OH} < 45^\circ$ (the angle roughly separating “upright” from “parallel”), the O and H atom velocities of the considered aluminol are velocity swapped, otherwise not. According to this method, not all (four, see Fig. 1) non-hydrogen bonded aluminols are usually excited at the beginning of phase (3).

B. Analysis of NE-AIMD results

In order to analyze the dynamics, two sets of calculations were carried out. In a first set, we analyzed the time-dependent, ensemble-average kinetic energies of atoms or atom groups after excitation. Analyzing the kinetic energy of the excited aluminol OH groups and fitting them to analytic functions, defines a vibrational relaxation time. Analyzing the kinetic energy content in other subsystems, such as hydrogen-bonded aluminols, the water layer, or the alumina surface, gives insight into the pathways of energy flow, *i.e.*, into the relaxation mechanism.

An alternative analysis method, is to employ “instantaneous kinetic energy spectral densities” for non-equilibrium MD, as proposed in ref. 27 and 48–50. These are obtained *via* Fourier transforming mass-weighted velocity–velocity autocorrelation

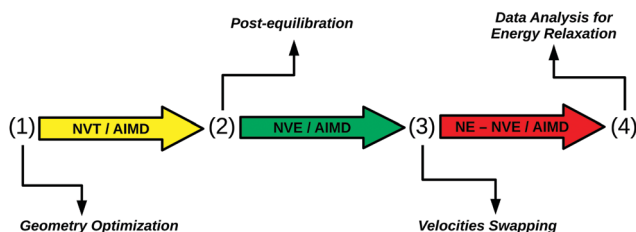


Fig. 2 Schematic representation of non-equilibrium AIMD protocol: in phase (1), first the initial HS + 2ML structure is optimized at the PBE + D2 level of theory. Then, canonical *ab initio* molecular dynamics (*NVT*/AIMD) trajectories at 300 K are propagated until thermal equilibrium is reached after 24 ps. At (2), AIMD trajectories are further propagated within the microcanonical ensemble to remove any spurious effect caused by coupling with the thermostat (*NVE*/AIMD). This phase results in a “post-equilibration”. After a certain time (10 ps for *NEQ*₁ and 20 ps for *NEQ*₂), nuclear velocities of specific O–H bonds are rescaled or “swapped”, at the beginning of phase (3). Then, a “non-equilibrium” phase begins during the course of which excited O–H kinetic energies are monitored, for 10 ps (*NE-NVE*/AIMD). Finally, in step (4) data are analyzed to unravel details of the vibrational relaxation process.

functions in individual time windows along the AIMD trajectories. Here we use a variant of this approach by considering the Fourier transforms of non-mass weighted VVAFs, by which we obtain Vibrational Density Of States (VDOS) curves as a function of both frequency ω and delay time Δt . The latter is an approximation to a time-dependent vibrational spectrum. We should point out that a similar procedure has been proposed by Lesnicki *et al.* to study vibrational relaxation of water, with special application for the liquid and the CaF₂/H₂O interface,^{51,52} but here the computed spectral densities are directly obtained by VVAFs. The analysis based on VDOS($\omega, \Delta t$) can also be done atomwise or for various subsystems of HS + 2ML. Specifically, the total VDOS($\omega, \Delta t$) is calculated from real parts of the Fourier transforms as

$$\text{VDOS}(\omega, \Delta t) \propto \int_{\Delta t-T}^{\Delta t} C_{\text{vw}}(t; \Delta t) \cos(\omega t) w(t) dt. \quad (3)$$

Here, Δt indicates the time delay since the beginning of phase (3) from Fig. 1, while T is the time interval over which the correlation function is computed at time Δt . We employ $T = 1$ ps in this work in order to balance an adequate spectral resolution with appropriate time sampling of AIMD trajectories. We also adopt a window function $w(t)$ for the Fourier transform, with $w(t) = \cos^2(\pi t/2T)$. Further, the VVAFs $C_{\text{vw}}(t; \Delta t)$, which indirectly depend on time Δt , are computed as

$$C_{\text{vw}}(t; \Delta t) = \frac{1}{N_t N} \sum_{i=1}^N \sum_{j=1}^{N_t} \langle v_i(t_j; \Delta t) v_i(t_j + t; \Delta t) \rangle. \quad (4)$$

Here, N_t is the number of timesteps employed to sample the correlation function in the interval $[\Delta t - T, \Delta t]$ and N is the total number of (moving) atoms. The time variable is $t_j = (j - 1)\delta t$, where $\delta t = 0.2$ fs as for all AIMD calculations here. Finally, $\langle \dots \rangle$ denotes an ensemble average (over all trajectories). Since each total VVAF is normalized for the given time interval $T = 1$ ps, their integral remains constant. In what follows, we evaluate VDOS($\omega, \Delta t$) during phase (3) of our AIMD trajectories, at various snapshot times Δt after “IR excitation”.

IV. Results and discussion

A. Time-dependent average kinetic energies and relaxation

In the following section results from the non-equilibrium AIMD trajectories are reported, for the HS + 2ML system shown in Fig. 1 with excited non-hydrogen bonded aluminols. We will only refer to trajectory-averaged properties in time intervals 10 ps before and 10 ps after “IR excitation” and quantities derived thereof. For this purpose, we average all ten trajectories sampled during this time and do not distinguish between NEQ₁ and NEQ₂ groups anymore. In fact, averaging separately over the two groups gives very similar results.

We first show in Fig. 3(a), the total kinetic energy $E_{\text{kin}}(t) = \frac{1}{2} \sum_i^N m_i v_i^2(t)$ for phases (2) and (3). In this figure, $t = 10$ ps refers to the instant of “IR excitation”. We note that the velocity swapping at $t = 10$ ps, leads to a total kinetic energy gain by

around 1.4 eV (from ~ 2.7 to ~ 4.1 eV), right after the excitation. Given $\hbar\omega_0 = 3750 \text{ cm}^{-1}$ (0.46 eV), this corresponds to the excitation of around three aluminols on average, with one quantum each. Since only sufficiently upright non-hydrogen bonded aluminols are excited, this reflects the fact that not all four nhb-upright aluminols found in the static structure of Fig. 1 are excited in average, due to their dynamical fluctuations.²³

A more detailed analysis of the time evolution of the kinetic energy before and after velocity swapping is provided in panels Fig. 3(b) and (c), for different subsystems of the HS + 2ML model. In particular, we report averaged E_{kin} values for four different subsystems:

- the four non-hydrogen bonded aluminols, among them the excited, “upright” OH bonds (“nhb-OH”);
- the hydrogen-bonded aluminols (“hb-OH”);
- the water molecules forming the adsorbed layer (“H₂O”);
- and the remaining surface atoms (“Al₂O₃”).

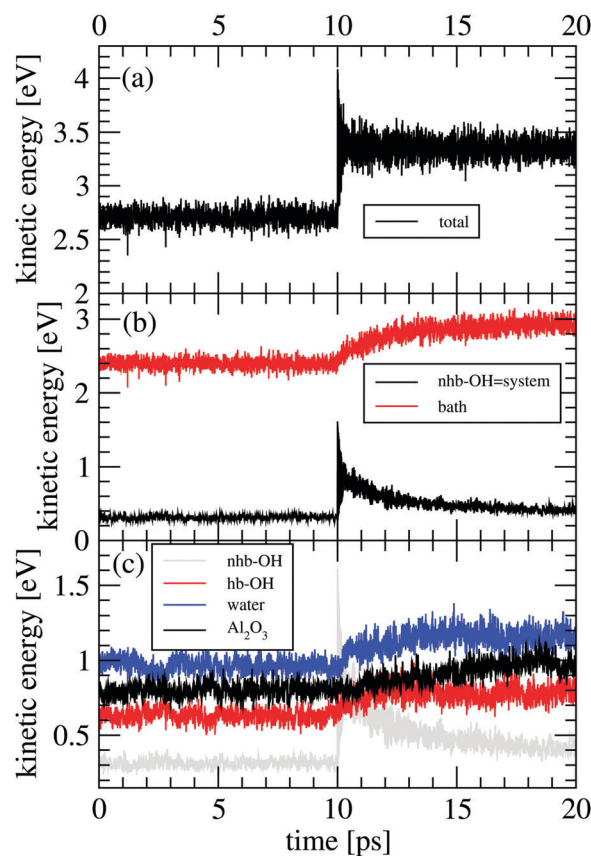


Fig. 3 Analysis of kinetic energies during NE-NVE/AIMD simulations. (a) Averaged (over ten trajectories) kinetic energy of the total HS + 2ML system. (c) Analysis of the averaged kinetic energy of different subsystems, “H₂O” (the water layer), “Al₂O₃” (the alumina surface), “hb-OH” (the hydrogen-bonded aluminol OH groups), “nhb-OH” (the non-hydrogen bonded aluminol OH groups). (b) Coarse-grained analysis of averaged kinetic energies, showing kinetic energies of the “system” containing the chromophores, $E_{\text{kin}}^{\text{sys}} = E_{\text{kin}}^{\text{nhb-OH}}$, and the “bath” consisting of the rest, $E_{\text{kin}}^{\text{bath}} = E_{\text{kin}}^{\text{hb-OH}} + E_{\text{kin}}^{\text{H}_2\text{O}} + E_{\text{kin}}^{\text{Al}_2\text{O}_3}$. In all cases, the time $t = 10$ ps corresponds to the time of “IR excitation”.

A still quite coarse-grained picture of the relaxation dynamics is provided in Fig. 3(b), showing the kinetic energy of a “system” containing the non-hydrogen bonded aluminols only, $E_{\text{kin}}^{\text{sys}} = E_{\text{kin}}^{\text{nhb-OH}}$, and a “bath”, $E_{\text{kin}}^{\text{bath}} = E_{\text{kin}}^{\text{hb-OH}} + E_{\text{kin}}^{\text{H}_2\text{O}} + E_{\text{kin}}^{\text{Al}_2\text{O}_3}$ comprising all the movable rest. The black curve in panel (b) clearly reflects a quasi-exponential decay for excited aluminols (the “system”), while the average “bath” kinetic energy (red) follows an incremental trend which reaches a new steady-state like condition within the last picoseconds of the non-equilibrium AIMD. From the “system” curve, we may extract a vibrational relaxation time, τ , for the non-hydrogen bonded aluminols, by fitting their average kinetic energy, $E_{\text{kin}}^{\text{nhb-OH}}$, for $t > t_0$, with an exponential curve:

$$E_{\text{kin}}^{\text{nhb-OH}}(t - t_0) = E_{\text{kin}}^{\text{nhb-OH}}(t_0)e^{-(t-t_0)/\tau} + \Delta E_{\text{kin}}^{\text{nhb-OH}} \quad (5)$$

Here, t_0 denotes the time of IR excitation of the chromophores. Further, $\Delta E_{\text{kin}}^{\text{nhb-OH}}$ represents the long-time limit of $E_{\text{kin}}^{\text{nhb-OH}}$. A best fit give $\tau = 2.0 \pm 0.37$ ps and $\Delta E_{\text{kin}}^{\text{nhb-OH}} = 0.42$ eV, respectively.

In Fig. 3(c), we further disentangle the kinetic energy flow from the “system” into various “bath” subsystems. From the three subsystem curves shown there, we note that both neighbouring water molecules ($E_{\text{kin}}^{\text{H}_2\text{O}}$) and hydrogen-bonded aluminols ($E_{\text{kin}}^{\text{hb-OH}}$) are quite effective channels for accepting vibrational energy from the non-hydrogen bonded aluminols: the average kinetic energies of the water layer and the hb-OH bonds increase rapidly within the first 2–3 ps after vibrational excitation of the chromophores, reaching a new plateau corresponding to a “hot ground state” after 5 ps. In contrast, surface atoms ($E_{\text{kin}}^{\text{Al}_2\text{O}_3}$) seem to increase their kinetic energy more slowly, time-delayed (after 5–6 ps), and to a lesser extent. Within the present model at least, a direct energy transfer from the excited aluminols to the alumina surface phonons is of

minor importance for the relaxation process, in contrast to more “lateral” transfer to neighbouring water molecules or aluminol groups. This is consistent with the mismatch of frequencies, with O–H vibrations being much more energetic than surface phonons. It is also in line with test AIMD calculations according to which excited aluminol O–H vibrations, within the present model at least, on a hydroxylated but otherwise water-free alumina surface, showed no signs of vibrational relaxation up to timescales of ten picoseconds whatsoever (not shown here).

B. Time-dependent vibrational densities of states (VDOS)

The second analysis tool mentioned earlier is the time-windowed VDOS according to eqn (3). In what follows, we evaluate $\text{VDOS}(\omega, \Delta t)$ during phase (3) of our AIMD trajectories, at ten snapshot times $\Delta t = 1, 2, \dots, 10$ ps after “IR excitation”. The VDOS curves are shown in Fig. 4. For comparison, we will also refer to VDOS curve obtained from equilibrium *NVT*/AIMD calculations (300 K) done in ref. 23, and a *NVE*/AIMD spectrum for $\Delta t = 0$, *i.e.*, from prior to the excitation step.

Looking first at the equilibrium situation, which is shown in Fig. 4(a), black curve, we find the narrow, “upright” nhb-aluminol peak at 3750 cm^{-1} mentioned earlier, while hydrogen-bonded aluminol and water–OH groups form a broad band from about 3000 to 3600 cm^{-1} . At around 1600 cm^{-1} we see the water bending modes and below $\sim 1200 \text{ cm}^{-1}$ water librational, OH bending, substrate phonons and water layer vibrations, all similar to those found from NMA. The assignment comes from NMA and also from atomic or subsystem analysis of the VDOS, as outlined in ref. 23. Note that the low-frequency band has the highest VDOS, followed by the higher-frequency regions.

Let us look now at the time evolution of $\text{VDOS}(\omega, \Delta t)$ in the non-equilibrium situation, shortly after “IR excitation” of selected nhb-aluminol chromophores, at $\Delta t = 1$ ps. The corresponding VDOS is also shown in Fig. 4(a), blue curve, and also in Fig. 4(b) when zoomed into the O–H stretching range $[3000, 4000] \text{ cm}^{-1}$. In

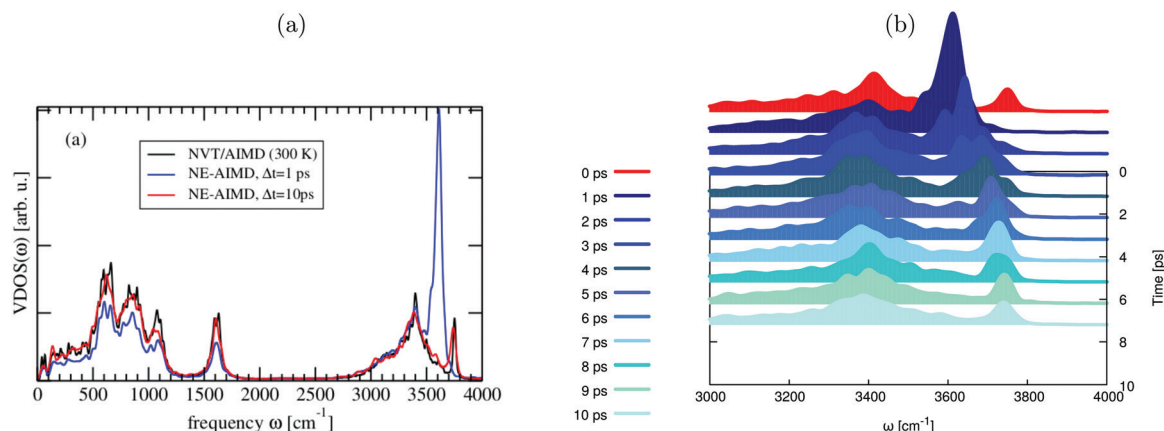


Fig. 4 Vibrational density of states, VDOS, for the total HS + 2ML system. (a) The VDOS from an equilibrium *NVT*/AIMD (300 K) calculation (black) in comparison to VDOS curves obtained from *NE-NVE*/AIMD, with time delays $\Delta t = 1$ ps (blue) and $\Delta t = 10$ ps (red), respectively, all in the full frequency range $[0, 4000] \text{ cm}^{-1}$. (b) VDOS spectra from *NE-NVE*/AIMD simulations at various delay times, in the O–H stretching region $[3000, 4000] \text{ cm}^{-1}$. Time is with respect to the beginning of phase (3), *i.e.*, after “IR excitation” in all cases. $\Delta t = 0$ in (b) refers to the situation right before excitation, giving a spectrum very similar to the *NVT*/AIMD (300 K) one.

this case, we note the very intense peak around 3600 cm^{-1} . This peak in the VDOS at $\Delta t = 1\text{ ps}$ arises from the vibrationally excited non-hydrogen bonded aluminols and is now most intense. Its position at a lower frequency compared to the room temperature equilibrium situation, not centered around 3750 cm^{-1} , can be explained by taking into account both the anharmonicity of the pre-excited O–H bond and the classical motion of nuclei.^{27,48} In contrast, at room temperature without O–H pre-excitation, the trajectories explore predominantly the harmonic, low-energy portion of the O–H potential curve.

With increasing delay time Δt after pre-excitation, we see from Fig. 4 that the 3600 cm^{-1} peak loses intensity, and is being blue-shifted towards a vibrational frequency of $\sim 3740\text{ cm}^{-1}$. In the classical dynamics picture, the blue-shift arises from the fact that O–H vibrations lose energy and amplitude, and the corresponding trajectories then also start to probe the harmonic low-energy regions of the O–H potential curves. The $\sim 3740\text{ cm}^{-1}$ signal is rather stable in position and intensity, after a few ps. Some, smaller modifications of VDOS curves can be also noticed for the hydrogen-bonded modes, located between 3000 and 3500 cm^{-1} , as seen in Fig. 4. Further, the distinct H_2O bending-mode peak around 1600 cm^{-1} and the low-frequency modes below 1100 cm^{-1} change in intensity while their position remains largely unaffected.

The VDOS spectrum at $\Delta t = 10\text{ ps}$ closely reflects all the features of the equilibrium 300 K spectrum, as can be most clearly seen from Fig. 4(a). This spectrum is also similar to the non-equilibrium, pre-excitation spectrum at $\Delta t = 0$, which has been added in Fig. 4(b) (in red). Remaining differences between $\text{VDOS}(\omega, \Delta t = 10\text{ ps})$ and the equilibrium 300 K spectrum can be due to the fact that shorter time-intervals have been used for Fourier transformation in the Δt -resolved case, and also since the system contains more energy and thus has a higher effective temperature after excitation. Altogether, the time-resolved spectrum reflects the approach to thermal equilibrium, following the excitation.

All of these findings are also qualitatively consistent with what has been found above for the relaxation process, from kinetic energy distributions. In order to directly address the spectral changes due to vibrational energy redistribution in time, we define the following function, $\Delta\text{VDOS}(\Delta t)$, by integrating for a given Δt over frequency, and referring to $\text{VDOS}(0)$, which is the VDOS sampled within the last 1 ps of the equilibrium-NVE phase (2), *i.e.*, right before velocity swapping:

$$\Delta\text{VDOS}(\Delta t) = \int_0^\infty [\text{VDOS}(\omega, \Delta t) - \text{VDOS}(\omega, 0)] d\omega. \quad (6)$$

Again, this quantity can be decomposed into atomic or subsystem contributions.

In Fig. 5, we report the integrated VDOS curves for different delay times, *i.e.*, $\Delta\text{VDOS}(\Delta t)$ in eqn (6). We employ again a constant time interval of 1 ps . The analysis is done for the total HS + 2ML system, but also for subsystems. In particular, we consider hydrogen- and non-hydrogen bonded aluminols (“hb-OH” and “nhb-OH”), their sum (“aluminols, (nhb + hb)-OH”), the water layer (“ H_2O ”), and the surface (“ Al_2O_3 ”).

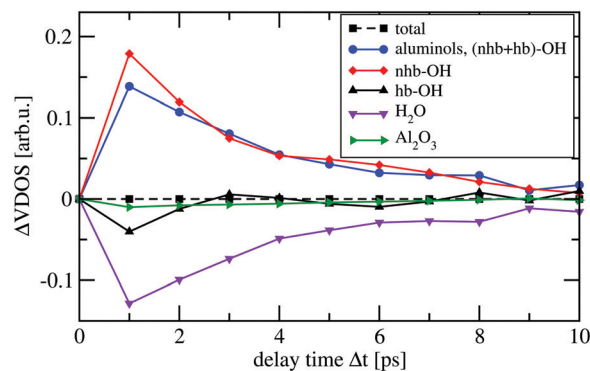


Fig. 5 Time-evolution of the integrated change of VDOS curves, $\Delta\text{VDOS}(\Delta t)$, from non-equilibrium AIMD simulations, averaged over all ten trajectories. The analysis is done for the total system and for various subsystems.

Since each total VVAF is normalized, the integrated total spectral density doesn't change as a function of time, so that the total $\Delta\text{VDOS}(t)$ remains constant at 0 (dashed black curve). On the other hand, spectral densities of individual subsystems manifest significant changes during the 10 ps of propagation. The $\Delta\text{VDOS}(\Delta t)$ of non-hydrogen bonded aluminols (red), suddenly increase between 0 (end of phase (2)) and 1 ps , then progressively decay by at least 85% within the 10 ps propagation time. As we would expect, the integrated spectral densities of hydrogen-bonded aluminols (blue), water layer (magenta) and surface atoms (in green) have an increasing behaviour within at least the very first 2 or 3 ps after 1 ps . This can be interpreted as a vibrational energy flow into these subsystems, which are all part of the “bath”. We also see that the water layer seems most affected, followed by the hydrogen-bonded aluminols and the surface, which shows only very small changes in $\Delta\text{VDOS}(\Delta t)$. While the inactivity of the surface atoms in relaxation is obvious, the relative importance of individual water molecules *vs.* hydrogen-bonded aluminols is debatable, because there are more water atoms than hydrogen-bonded aluminols. At longer times, for the various bath subsystems, we see oscillatory behaviours. These are probably due to energy exchange within different dissipating channels, but also finite size effects cannot be excluded.

Due to the relative inactivity of the surface atoms, the blue ΔVDOS curve, which is associated with both non-hydrogen bonded and hydrogen-bonded aluminols, is close to a mirror image of the integrated spectral density of all water molecules (magenta). This reinforces the hypothesis that vibrational energy relaxation happens by coupling to modes of energy between 3000 and 3500 cm^{-1} , which are located in the hydrogen-bonded VDOS band. Further, when looking at the black solid and the magenta ΔVDOS curves, we also notice that spectral densities of hydrogen-bonded aluminols and water molecules evolve almost symmetrically, which may indicate an efficient energy exchange between these two, strongly hydrogen-bonded subsystems.

Finally, by inspection of Fig. 5, we could also deduce that the main relaxation happens within the first 4 ps or so, and shows signatures of non-exponential behaviour. This timescale is

somewhat slower than the one derived from kinetic energy analysis (~ 2 ps). In passing, we note that preliminary experimental results for this system suggest indeed O–H relaxation times in the order of a few picoseconds.²⁶ All of this indicates that the present situation with an O–H chromophore which is relatively weakly coupled to a surface and water environment, will lead to lifetimes between “short” (sub-ps) and “long” (many ps or even longer), the latter two for strongly and very weakly coupled systems, respectively.

V. Summary and conclusions

To summarize, the analysis of both kinetic energy and vibrational state densities provides qualitative and partially quantitative tools to unravel the energy redistribution of excited upright non-hydrogen bonded aluminols, which were the focus of this work. Our calculations suggest that the latter subsystem of the HS + 2ML surface is characterized by a vibrational band located between 3700 and 3800 cm^{-1} , outside the other high-frequency bands which are associated with hydrogen-bonded modes. The analysis of non-equilibrium AIMD trajectories shows that, once the non-hydrogen bonded, upright aluminols are excited, they relax rapidly within the first picoseconds of propagation. Relaxation times obtained either *via* an exponential fit of their average kinetic energy, or by looking at the time-evolution of the frequency-integrated spectral densities, both suggest a “lifetime” between 2 and 4 ps. Compared to other studies reporting on O–H vibrational dynamics, especially the analysis of Borguet *et al.*,^{15–17} our findings indicate an intermediate relaxation timescale for an O–H chromophore with weak coupling to its aqueous environment. Our tools also allow us to disentangle the role of various other subsystems of the HS + 2ML model for the relaxation process. Indeed, we can identify as main dissipative channels both hydrogen-bonded aluminols and, in their sum even more important, water molecules in the adsorbed water layer. In general, hydrogen-bonded modes with frequencies between 3000 and 3500 cm^{-1} act as efficient pathways for the redistribution of the initial excess of kinetic energy in quasi-free, high-frequency O–H aluminols.

Methodologically, here we combined non-equilibrium AIMD methods with time-dependent correlation function techniques. The vibrational excitation is simulated by employing a “velocities swapping” method, which allows for a mode-selective perturbation by adding a given excess of kinetic energy, *i.e.* a vibrational quantum in our approach, all within the framework of classical dynamics. The relatively high increase of kinetic/thermal energy, accompanied with the inherent structural fluctuations of AIMD trajectories, while causing on one hand an overall heating of the system, higher than in the actual experiments, can also account for multi-photon excitations as the ones due to spectrally-broad exciting IR pulses.

On a more general level, it shall be underlined that while the AIMD approach is “costly”, it has the advantage of also allowing to treat reactive processes, and it is completely general, *i.e.*, not restricted to a system class for which a force field, say, has been

parametrized. Further, the NE-AIMD/TCF methodology opens the way to real, time-resolved vibrational spectra, *e.g.*, time-resolved IR or VSF spectra. In this case, dipole–dipole auto- or dipole-polarizability cross-correlation functions need to be determined, which can also be done with the help of velocity–velocity autocorrelation functions.^{22,23,45} Work along these lines is in progress in our laboratory.

Conflicts of interest

There are no conflicts to declare.

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