1. Introduction

The stabilities, structures, and reactivities of metal oxide surfaces and nanoparticles under ambient conditions are strongly affected by their hydration-hydroxylation. Here especially the water molecules residing closest to the metal oxide are targets of intense scientific interest as they are the molecules most drastically modified by the surface and often even subject to dissociation. Spectroscopic signals from dissociated and intact water molecules on many metal oxide surfaces have been presented in the literature over the years (see, for example, ref. 1–9) and here vibrational and electron spectroscopies have emerged as being especially informative. Thus, recently, some of the seemingly most simple and prototypic water/oxide systems, such as H$_2$O/MgO(100), H$_2$O/CaO(100), H$_2$O/ZnO(1010) and H$_2$O/rutile-TiO$_2$(110), have been targets of vibrational spectroscopic studies of various flavors such as IRRAS (Infrared reflection-absorption spectroscopy), SFG (sum-frequency generation), and FTIR (Fourier transform Infra-red) spectroscopy with an ATR (attenuated internal total reflection) device.

Theoretical investigations are critically important in the field of vibrational spectroscopy, and most of the experimental studies today contain supporting vibrational calculations.

The special value of calculations is that they can provide both detailed local structure information and detailed, well resolved, spectral information: together these two advantages help to decipher complex experimental spectra for complex systems. Moreover, such calculations often have the power to take us further and provide an explanation, or a rationale, behind the experimentally observed spectral features. The objective of this paper is to present a unifying, systematizing framework that relates the vibrational dynamics of surface OH groups to their chemical environment for both OH$^-$ and H$_2$O simultaneously.

We will show that in the analysis of experimental or calculated OH spectra it is not helpful to treat all groups as water-like. We find that all the dissociated OH groups present for different water coverages on MgO(001) and CaO(001) are indeed hydroxide ions, and this has consequences for how they respond to perturbations from their environments. Thus we find that the intact water molecules on the surface are always downshifted with respect to the unperturbed (gas-phase) molecule while the OH$^-$ groups are either upshifted or downshifted with respect to the gas-phase ion. We will also show that in spite of these differences it is possible to treat the surface water molecules and surface hydroxide ions on an equal footing, using a simple and intuitive electrostatic analysis involving the electric field from the surroundings as the descriptor. The electric field has been found to be quite a useful local useful local descriptor in the context of the OH vibrational downshift of perturbed water molecules in various situations, such as in uniform electric fields in the gas phase, and H$_2$O simultaneously.
liquid water,14,15 and in crystalline hydrates.16 However, as far as we are aware, our treatment in this paper represents a new way of looking at OH vibrations on ionic surfaces. Our hope is that our new model will simplify future interpretations of water OH vibrational spectra for surface systems where electrostatic interactions play an important role, which is the case on metal oxide surfaces.

2. Systems and methods

2.1 Systems

Water clusters (monomers, dimers, and trimers) on MgO(001) and CaO(001) as well as overlayers of 1.0 and 1.25 monolayer (ML) coverages are studied in this paper. Two of these structures are shown in Fig. 1. A full monolayer is defined as 1 adsorbed water molecule per surface cation (or formula unit). The starting geometries for the geometry optimizations were taken from the previously reported 1.0 ML and 1.25 ML over-layer structures on MgO(001) and CaO(001) by Włodarczyk et al.2 and Zhao et al.,5 respectively. For the overlayers and the trimer on MgO(001), some of the water molecules dissociate and form two types of hydroxides, namely O\(_h\)H or OH\(_f\). The s in O\(_s\) stands for “surface” and the f in H\(_f\) stands for “free”. Due to the stronger basicity of CaO, water molecules near its surface are more likely to dissociate than in the case of MgO, and to give rise to many non-equivalent OH groups.

Altogether we examine 56 structurally unique intact H\(_2\)O molecules and 34 structurally unique OH\(^-\) ions, covering a range of different situations from weak to strong “intermolecular” interactions. The H\(_2\)O/MgO(001) and H\(_2\)O/CaO(001) systems were modelled as 3-D periodic boxes, consisting of slabs of four substrate layers, plus the water adsorbate layer and a 15 Å thick vacuum region. The calculated lattice constants of 4.25 Å for bulk MgO(001) and 4.83 Å for bulk CaO(001) were used to build the clean slabs. In all interface calculations, water molecules were adsorbed only on one side of the slab.

Gas-phase systems. The isolated gas-phase water molecule and hydroxide ion were optimized with 3D periodic boundary conditions in 14 Å × 15 Å × 16 Å simulation boxes.

![Fig. 1](image-url) Top and side views of two of the many interface structures examined in this study. (a) H\(_2\)O monomer/MgO(100), (b) 1 ML H\(_2\)O/MgO(100), with its three inequivalent water molecules w1, w2 and w3, where w1 is dissociated and w2 and w3 intact. The cartoon in (c) defines the three types of surface OH groups. The colour scheme is: yellow atoms = Mg, burgundy red atoms = O (surface lattice oxide), light red atoms = O (intact adsorbed water molecules), orange atoms = O (OH\(_s\)), white atoms = H.

2.2 Electronic structure calculations

We recently found17 that the optPBE-vdW functional18 predicted the adsorption energy of water on MgO(001) in agreement with the experimental isosteric heat of adsorption, and we have chosen this functional for our study here. All calculations were performed with the VASP program.19-22 The electron-core interactions were described using the PAW formalism.23,24 For H, O, Mg and Ca, the 1s\(^1\), 2s\(^2\)p\(^1\), 2p\(^3\)s\(^2\) and 3p\(^5\)s\(^2\) electrons, respectively, were treated as valence electrons. All calculations were performed with a plane-wave kinetic energy cutoff of 400 eV and a 4 × 4 × 1 Monkhost–Pack k-point mesh. Only the \(\Gamma\) point was used for the gas-phase water molecule and hydroxide ion. We used Gaussian smearing with a width of 0.1 eV in all cases.

Full geometry optimizations were performed for all atoms in all systems, except that the atoms residing in the two “bottom” layers of the ionic slabs were kept fixed. The conjugate-gradient algorithm was used for the structure optimizations and cell parameters and atomic coordinates were considered to be converged when the forces acting on the atoms were less than 0.0021 eV Å\(^{-1}\). A convergence criterion of 10\(^{-7}\) eV for the total energy was used in the electronic self-consistent calculations.

2.3 Calculation of anharmonic OH frequencies

We have calculated all OH stretching vibrational frequencies using a 1-dimensional (1D) uncoupled OH vibrational model. In this model, each OH potential energy curve is constructed so as to allow the vibrating OH bond (in a H\(_2\)O molecule or OH\(^-\) group) to contract and stretch along the OH bond, keeping the center of mass of the OH group fixed while the remaining parts of the system were kept fixed at their optimized positions. We used a reduced mass of 0.94808 amu for the vibrating OH oscillator and 19 energy points were generated along the potential energy curve with a step size of 0.06 Å. More precisely, we used 12 energy points above and 7 below the equilibrium OH distance, \(r_c\).

The 1D vibrational Schrödinger equation was then solved for the vibrational energy levels using the discrete variable basis-set representation (DVR) approach of Light et al.25,26 The anharmonic stretching frequencies were calculated by taking the energy difference between the ground and first excited vibrational eigenstates.

This computational approach has previously been used by us to calculate OH frequencies of water molecules and hydroxide ions in crystals16,27,28 and on surfaces29 as well as in liquid water30 and ionic aqueous solutions.31

2.4 Calculation of electric field from the environment

The external electric field exerted on the surface OH groups from its surroundings is a central concept in this study. For an OH oscillator that belongs to an intact water molecule all neighbours outside that water molecule constitute the external world. Other appropriate names for this field might have been the \textit{in situ} field, on-site field, or perhaps intrinsic field to pinpoint that we are not dealing with any externally applied...
electric field, but we have chosen the term external field to emphasize that it is generated by all the short- and long-range neighbours outside the targeted hydroxide ion or water molecule.

Wannier orbitals and their centres\textsuperscript{32} were calculated from the final electronic wave function using the WANNIER90 program.\textsuperscript{33} The electric field generated by the surroundings at a probe site was calculated from the positions and charges of the Wannier centres and the nuclei (with any accompanying core electrons as described by the PAW pseudopotentials) using the GULP\textsuperscript{34} program. The on-site electric field calculated with this program contains both the self-field of the probed water molecule (or hydroxide ion) itself and the external electric field from the surroundings. Since we want to describe the effect of the gas-to-surface OH frequency shifts in terms of the surroundings only, the self-field generated by the internal charge sites of the probed water molecule or hydroxide ion were removed “manually” from the total on-site field value.

For each investigated OH group, in each of our structurally optimized surface systems, the equilibrium position of the H nucleus was selected as the probe site where the electric field strength was evaluated, and it is the fragment component along the OH bond that was monitored, as we have found that the electric field components perpendicular to the vibrating OH bond have only a negligible effect on the OH frequency. We denote it $E_i$, sometimes $E_i@H$. The field direction is defined according to the standard physics definition (i.e. from plus to minus).

2.5 Calculation of molecular dipole moments and dipole moment derivatives

The dipole moments of the isolated ($\mu^0$) water molecule and hydroxide ion were calculated from the positions and charges of their nuclei and four occupied Wannier centers, and similarly for the species on the surface ($\mu^{\text{bound}}$). The induced dipole moment was calculated from $\mu^{\text{bound}} - \mu^0$. The dipole moments were calculated with respect to the center-of-mass of the oscillating OH bond. For water, the dipole moment value does not depend on the reference position. For the charged hydroxide ion it does, but the expressions that we will make use of involve the first derivatives of dipole moments with respect to $r_{\text{OH}}$, and therefore the choice of reference point will again be unimportant. The dipole moment derivative was evaluated at the bound state equilibrium distance $r_{\text{e,bound}}$, using a 5-point numerical differential formula.

3. Results

3.1 The electric field as the origin of the gas-to-surface frequency shifts

The calculated uncoupled, anharmonic OH frequencies, $\nu_i$(OH), for the intact water molecules, adsorbed on either MgO(001) or CaO(001) are plotted in Fig. 2a. They span a large range, from about $3600 \text{ cm}^{-1}$ down to about $2100 \text{ cm}^{-1}$. Actually, this is also true for most of the individual adsorbate systems in the figure, except for the smallest water clusters. Note that the spread is not due to couplings, as these are uncoupled vibrations; instead, the spread directly mirrors the spread in the strengths of the external perturbations exerted from the surroundings on the individual water molecules. Going new to the dissociated water molecules in Fig. 2c, we find somewhat more structure in the data points: one highly populated group of frequencies between about $3600$ and $3470 \text{ cm}^{-1}$, immediately below which there is a more sparsely distributed group with values all the way down to about $2000 \text{ cm}^{-1}$. We can identify the uppermost group of frequencies with the structural motif that is commonly referred to as the OH$\cdot$ type in the literature. The frequencies lying about $3410$ and $3300 \text{ cm}^{-1}$ originate from O-H groups that are weakly bound to OH$\cdot$. Finally, the more low-lying set of frequencies also originate from O-H groups, but those which are strongly bound to the OH$\cdot$. In order to bring some logics and systematics into the presentation of all these data we display the same data in Fig. 2b and d, but now plotted against the probed electric field strength $E_i$. The result is impressive: all points nicely follow arc-like curves. The two curves are similar but not identical. The largest frequency in Fig. 2b lies close to $E_i = 0.0 \text{ a.u.}$, but in Fig. 2d it occurs at $E_i \approx 0.05 \text{ a.u.}$, below which no frequencies can be found since none of the hydroxide groups in our sample experiences small external fields. The latter may partly be a consequence of the relatively small tilts of the OH$\cdot$ groups from the surface normal compared to the water molecules which often lie flat on the surface (cf. Fig. 1b). It may also be so that environments giving smaller fields than 0.05 a.u. do not lead to water dissociation. In each of the two figures we have marked the frequencies at $E_i = 0.0$, i.e. the gas-phase frequencies. We note that for the intact water molecules all the frequencies are downshifted (red-shifted) with respect to this point while for the dissociated water molecules there exist oscillators which are upshifted (blue-shifted) with respect to the gas-phase value.

The two frequency vs. field curves just discussed are shown together in Fig. 3, together with the corresponding curves (dashed) obtained from MP2 calculations presented in ref. 10 for a water molecule and a hydroxide ion exposed to uniform electric fields of varying strengths directed along the OH stretch coordinate. The frequency-field curves in ref. 10 were found to be parabola-like for both water and hydroxide, but with their respective maxima at different positions. For H$_2$O the maximum was found\textsuperscript{35,36} to lie to the left of the zero-field line (cf. Fig. 3) resulting in redshifts for all positive (i.e. natural, realistic) fields. This is indeed consistent with experimental data: we are not aware of any experimental situation or system where an OH oscillator belonging to a bound water molecule is not red-shifted compared to the gas-phase water frequency. For the OH$^{-}$ ion in a uniform electric field, on the other hand, the maximum was found to lie on the positive field side.\textsuperscript{10} This implies that at small and intermediate fields, a frequency upshift with respect to the gas-phase frequency will occur, while with increasing fields finally very large downshifts appear. Indeed it is known from experiments, e.g. by Lutz and his co-workers,\textsuperscript{35-37} that hydroxide ions in crystalline hydroxides display both downshifts and upshifts compared to the gas-phase OH$^{-}$ frequency.

In this paper, we observe that the surface OH groups found on the hydrated-hydroxylated MgO and CaO surfaces qualitatively
follow the behaviour of the two ideal uniform-field systems. Thus, the intact surface water molecules essentially follow the uniform-field curve for H$_2$O, while the OH$_f$ (blue half-filled rings) and O$_s$H (red half-filled rings) data points display a similar behaviour to the uniform-field curve for the OH$^-$ ion. Bader-type atomic charges were calculated, and the net charges of the dissociated water OH groups in the various MgO and CaO surface systems studied here were found to lie in the range from $\sim 0.7$ to $\sim 0.9$ $e$, i.e. they can be classified as hydroxide ions. It is interesting to note that all the OH$_f$ groups in our sample accumulates close to the turning point of the frequency vs. field curve, while the O$_s$H groups lie further to the right on the curve. The former are all blueshifted with respect to the gas-phase OH$^-$ ion, while all of the O$_s$H vibrations (except one) are redshifted, most of them even strongly so. We believe that these findings are likely to hold for hydroxylated metal oxide surfaces in general, at least in the absence of a second water layer.

3.2 Perturbation from the surface and the co-adsorbates

In this section, we will provide a rationale behind the different water and hydroxide OH frequency redshift and blueshift...
behaviour displayed in Fig. 3. One of our intact surface water molecules is used as the example in Fig. 4a, namely the water monomer on the MgO(001) surface, and more specifically the OH leg which binds to the surface (cf. Fig. 1a). The black curve in the figure ($V_{\text{gas}}$, or equivalently, $V_0$) is the calculated potential energy for the isolated gas-phase molecule, with $r_{\text{OH}}$ being the uncoupled OH stretching coordinate. The red curve marked $V_{\text{bound}}$ is the corresponding calculated potential energy for our selected water molecule on the surface, shifted so as to coincide with the unperturbed curve at $r_{\text{OH}} = 0.80$ Å.

The total perturbation exerted by the environment will be denoted the external potential $V_{\text{ext}}(r_{\text{OH}})$, defined by $V_{\text{ext}}(r_{\text{OH}}) = V_0(r_{\text{OH}}) + V_{\text{ext}}(r_{\text{OH}})$. $V_{\text{ext}}$ is shown in the upper half of Fig. 4a, again shifted so as to coincide with the other two curves at $r_{\text{OH}} = 0.80$ Å.

The $V_{\text{ext}}$ curve is seen to have a negative slope at the gas-phase equilibrium distance $r_{\text{OH}}$, and in fact everywhere in the figure, and we claim that the sign of the slope determines the sign of the gas-to-surface frequency shift; this claim will be substantiated in the coming sections.

We now turn to a dissociated water molecule. The bottom part of Fig. 4c presents three parabola-like potential energy curves, one for a surface-bound $O_s\text{H}$ hydroxide ion (red curve), one for a gas-phase hydroxide ion (black curve) and one for a surface-bound OH$_f$ hydroxide ion (blue curve). The upper part of the figure shows the corresponding $V_{\text{ext}}$ potential energy curves. As in the previous case, all curves were drawn to coincide for $r_{\text{OH}}$ equal to 0.8 Å. The chosen O$_s$H ion is one of the strongly red-shifted OH$^-$ groups in Fig. 3, and is seen to be accompanied by a $V_{\text{ext}}$ curve with a negative slope everywhere, while the OH$_f$ ion is accompanied by a $V_{\text{ext}}$ curve with a positive slope in Fig. 4c and it is one of the blueshifted OH groups. This means that our findings are in agreement with our claim above, i.e. that the sign of the slope is consistent with the direction of the frequency shift.

Ten more examples from our bound intact surface water molecules on MgO(100) and CaO(100) are shown in Fig. 4b and seven more of the bound hydroxides in Fig. 4d; a variety of strongly and weakly bound environments are represented in both cases. The water molecules are seen to all have $V_{\text{ext}}$ curves with negative slopes everywhere, while the hydroxide ions display either negative (always O$_s$H ions) or positive slopes (always OH$_f$ ions), and this is indeed consistent with our results in Fig. 2 and 3 where the O$_s$H groups were all found to be downshifted in frequency compared to the gas-phase ion, and the OH$_f$ groups were upshifted. The colour of each curve indicates the direction of our calculated gas-to-surface frequency shift.

In conclusion, Fig. 3 and 4 together paint a consistent picture. All blueshifted OH groups are exposed to an external...
perturbation with a positive slope, and all redshifted OH groups are exposed to an external perturbation with a negative slope. The effect of an external potential with a negative slope is to move the equilibrium distance to larger values (weaken the bond), and to make the potential energy curve flatter and the harmonic force constant smaller. The opposite in all respects is true for the blueshifted OH oscillators. The curvature of $V^{\text{ext}}$ never opposes the shift direction predicted by the slope.

For future reference, we now introduce some notations/definitions:

\[ V^{\text{ext}}(r_{\text{OH}}) = k^0_{\text{OH}}(r_{\text{OH}} - r_{\text{e,OH}})^2 + k^0_{\text{r}}(r_{\text{OH}} - r_{\text{e,r}})^3 + \ldots \]  
\[ \nu^{\text{bound}} = k^2_{\text{bound}}(r_{\text{OH}} - r_{\text{e,bound}})^2 + \nu^{\text{OH}}(r_{\text{OH}} - r_{\text{e,OH}})^3 + \ldots \]  
\[ V^{\text{ext}}(r_{\text{OH}}) = k^2_{\text{bound}}(r_{\text{OH}} - r_{\text{e,bound}})^2 + V^{\text{ext}}_{\nu}(r_{\text{OH}} - r_{\text{e,OH}})^3 + \ldots \]  

The centers of the expansions for the free and bound potentials are their respective equilibrium distances, thereby removing a linear term. For the external potential the center has been chosen to be $r_{\text{e,OH}}$. With these definitions we find that the shift of the harmonic force constant up to the quadratic term becomes:

\[ \Delta k_2 = k^2_{\text{bound}} - k^2_{\text{OH}} \approx \frac{3k^0_{\text{bound}}}{2k^0_{\text{OH}}} + k^2_{\text{OH}}. \]  

In a perturbative treatment both the harmonic frequency shift ($\Delta \nu = \nu^{\text{OH}} - \nu^{\text{bound}}$) and the anharmonic frequency shift ($\Delta \nu = \nu^{\text{bound}} - \nu^{\text{OH}}$) are related to $\Delta k_2$ through the following relations:

\[ \Delta \nu \approx \frac{\Delta k_2}{2k^0_{\text{OH}}} \]  
\[ \Delta \nu \approx \left[ (\nu^{\text{OH}} - \nu^{\text{bound}}) \right] \frac{\Delta k_2}{2k^0_{\text{OH}}} \]  

A derivation of the last two equations is presented in the Appendix.

Eqn (2) allows us to make a more quantitative comparison of the impacts of the $k^2_{\text{OH}}$ and $k^2_{\text{bound}}$ terms on the harmonic force constant $\Delta k_2$ and thereby also on the OH frequency shifts by way of eqn (3a) and (3b). Thus for each OH oscillator in our study, a quadratic fit to the VASP-calculated external potential $V^{\text{ext}}$ was performed and Fig. 5 displays $\Delta k_2$ as a function of the field at each OH oscillator. It is seen that for both intact and dissociated water molecules, eqn (2) is dominated by the $k^2_{\text{OH}}$ term. For fields up to about 0.1 a.u., which is the field range where the qualitative redshift/blueshift difference between water and OH$^-$ manifests itself, it is sufficient to approximate the external potential as being linear in the stretch coordinate. For both water and hydroxide, $k^2_{\text{OH}}$ is positive and $k^2_{\text{bound}}$ is negative, and so the sign of the prefactor of $k^2_{\text{OH}}$ is positive and the sign of $k^2_{\text{bound}}$ will determine the sign of $\Delta k_2$.

In the next subsection we present a simple electrostatic model to understand the rationale behind the field variations of the frequencies of the surface OH oscillators. In that model we will use eqn (2), setting $k^2_{\text{bound}} = 0$.

### 3.3 Interpretation – a dipolar model gives the direction

The number written to the right of each $V^{\text{ext}}$ curve in Fig. 4b and d is the field value that we have discussed extensively already, i.e. the field parallel to the vibrating OH bond, produced by surrounding charges, and evaluated at the bound H equilibrium position, $r_{\text{e,OH}}$. This is now our descriptor of the external environment of each surface OH group.

In ref. 10 and 11, as mentioned, isolated water and OH$^-$ species in uniform electric fields were studied, and it was shown that, in both cases, $V^{\text{ext}}$ is well approximated by a dipolar electrostatic model. Here we propose to use the same model with the uniform field being replaced by $E$. Using the expressions given in ref. 10 and 11 and applying eqn (1c), we get

\[ k^2_{\text{OH}}(r_{\text{OH}}, E) \approx -\left[ \frac{\partial \mu_{\text{OH}}}{\partial r_{\text{OH}}} \right] \frac{1}{E} - \int_{E_0}^{E_1} \left[ \frac{\partial \mu_{\text{OH}}}{\partial E} \right] \frac{1}{E} \, dE, \]

![Fig 5](image-url)  
Comparison of the VASP-calculated $\Delta k_2 = k^2_{\text{bound}} - k^2_{\text{OH}}$ values (black circles) with expression (2) in the paper, i.e. $\Delta k_2 = \Delta k_{\text{OH}}^{\text{to-surf}} \approx -[3k^2_{\text{OH}}/2k^2_{\text{OH}}] + k^2_{\text{OH}}$. Plotted against $E$. The red circles denote the $\sim [3k^2_{\text{OH}}/2k^2_{\text{OH}}]k^2_{\text{OH}}$ term and the blue circles denote the sum of the two terms. (a) The intact water molecules on MgO(001) and CaO(001). (b) The hydroxide groups on MgO(001) and CaO(001).
Here $\mu_0^{\text{OH}}$ is the electric dipole moment of the gas-phase (i.e. zero field) OH group when the OH stretch coordinate has the value $r_{\text{OH}}$, and $\mu_{\text{ind}}^{\text{OH}}(E_i)$ is the induced dipole moment. In accordance with eqn (1c), the derivative is taken at the gas-phase equilibrium distance. Since the field is taken at the equilibrium position of the bound hydrogen, also the dipoles must be calculated with respect to this point. However, since both the equilibrium position of the hydrogen atom and the center of mass of the OH groups do not change when $r_{\text{OH}}$ is changed, we may just as well use the latter as our reference point for calculating dipoles.

As we are primarily interested in a qualitative analysis, we assume that $d\mu_{\text{ind}}^{\text{OH}}(E_i)/dr_{\text{OH}}$ is approximately proportional to the field over an appreciably large range of field values. Eqn (4) then becomes

$$k_{\text{ext}}^{\text{OH}}(r_{\text{OH}}, E_i) \approx -d\mu_{\text{ind}}^{\text{OH}}/dr_{\text{OH}} \cdot E_i - \frac{1}{2} d^2\mu_{\text{ind}}^{\text{OH}}(r_{\text{OH}}, E_i)/dr_{\text{OH}}^2 \cdot E_i$$

(5a)

$$\approx -\mu_0^{\text{OH}} \cdot E_i - \frac{1}{2} \mu_{\text{ind}}^{\text{OH}} \cdot E_i^2$$

(5b)

The derivatives are displayed in Fig. 6. The symbol $x$ represents the $x_1$ component of the dipole polarizability tensor of the OH oscillator, and the prime denotes a differentiation with respect to the stretch coordinate, taken at the gas-phase equilibrium point $r_{\text{OH}}$. The $x'$ values were obtained from linear fits to our calculated $d\mu_{\text{ind}}^{\text{OH}}/dr_{\text{OH}}$ vs. electric field data, for small to modest field strengths.

Using eqn (2), (3) and (5) we then calculated the harmonic and anharmonic frequency shifts within our dipolar model and plotted the anharmonic shift in Fig. 7, together with the results from our VASP calculations. Given the many of approximations, the results are surprisingly good. We therefore accept the model as a reasonable approximation to reality and analyze the difference in behavior between the water and the hydroxide oscillators.

Inspection of eqn (5a) shows that the sign of $d\mu_{\text{ind}}^{\text{OH}}/dr$ and the relative magnitudes of $d\mu_0^{\text{OH}}/dr$ and $d^2\mu_{\text{ind}}^{\text{OH}}/dr^2$ determine the magnitude and sign of $k_{\text{ext}}^{\text{OH}}$, which in turn will determine the sign and magnitude of $\Delta\nu$ and the position of the maximum in the frequency vs. field correlation curves in Fig. 3. We find that all quantities in the second term of eqn (5a) are positive for all the surface OH oscillators while $d\mu_0^{\text{OH}}/dr_{\text{OH}}$ has different signs for water and hydroxide: $+0.62$ D Å$^{-1}$ for water and $-1.14$ D Å$^{-1}$ for hydroxide with our chosen DFT functional. It is the sign of $d\mu_0^{\text{OH}}/dr$ which is the ultimate origin behind the different frequency vs. field behavior of the water molecule and the OH$^-$ groups in Fig. 3. The negative sign for OH$^-$ will give rise to a parabola with a maximum for a positive field value.

This can easily be formalized as follows. From eqn (5b) we find that the frequency shifts as a function of the field lie on concave (since $x''$ is positive in all cases) parabola. The maxima of these parabolas occur where $E_i = -\mu_0^{\text{OH}} / x''$. Since $\mu_0^{\text{OH}}$ is positive for the water oscillator the maximum for the water parabola falls at a negative field value; a negative field value never occurs in the $in\ situ$ situations studied here, and seldom, if ever, in any equilibrium scenarios in nature either, as they would be energetically unfavourable. As a result the maximum occurs at zero field and all frequencies are downshifted. By a similar reasoning, since $\mu_0^{\text{OH}}$ is negative for the hydroxide oscillator, its maximum frequency shift occurs at a positive field value. The value of $k_{\text{ext}}^{\text{OH}}$ at this point is equal to $(\mu_0^{\text{OH}})^2 / 2x''$, which is positive. For field values between zero and $-2\mu_0^{\text{OH}} / x'' > 0$, the hydroxide oscillator will therefore be upshifted. In our present study no fields occur below $-2\mu_0^{\text{OH}} / x''$, and so all oscillators in fields between $-\mu_0^{\text{OH}} / x''$ and $-2\mu_0^{\text{OH}} / x''$ are upshifted.

The underlying reason for the different signs of $d\mu_0^{\text{OH}}/dr_{\text{OH}}$ for gas-phase water and hydroxide is related to the fact that when the gas-phase hydroxide ion is stretched, a very complex electron redistribution occurs, which leads to a net decrease in the dipole moment, and simultaneously makes it favourable.
for the hydroxide to contract when placed in an electric field. The water molecule, on the other hand, behaves like most molecules: the dipole moment increases if the OH bonds are stretched, and the system gains energy by elongating when exposed to an electric field.39

Further analysis performed by us suggests that, seemingly, the almost perfect agreement between the VASP and model results for water at high fields is at least partly due to error compensation. The neglect of the \( k_{2\text{ext}}^n \) term in the dipolar model is compensated by a too large slope of the \( k_{1\text{ext}}^o \) term in the same model, which in turn is likely due to the neglect of the non-electrostatic interactions (orbital overlap). For the hydroxide on the other hand, we find that the \( k_{2\text{ext}}^n \) term from the dipolar model is quite similar to that from VASP, and that it is mainly the lack of the \( k_{2\text{ext}}^o \) term that causes the discrepancy with the VASP results.

4. Summary and concluding remarks

This paper is an effort to rationalize the sign and size of the OH frequency shifts observed experimentally for water on metal oxide surfaces, namely \( \text{O}_2\text{H}\), \( \text{OH}_2\), and \( \text{OH}_w \) groups, the latter from intact water molecules. In total the paper analyses 90 unique surface OH groups – intact water molecules as well as hydroxide groups. The first part of the title of this paper is “Surfaces with both molecularly and dissociatively adsorbed water molecules and the OH groups are part of the same general scheme where the main behaviour of gas-to-surface OH frequency shifts on metal oxide surfaces”, which reflects the phenomenon that OHw groups as hydroxide groups. The first part of the title of this paper is unique surface OH groups – intact water molecules as well from intact water molecules. In total the paper analyses 90

The lack of the \( k_{1\text{ext}}^o \) term that causes the discrepancy with the VASP results.

Appendix: perturbation model

Since we want to discuss both the harmonic and the anharmonic frequency shifts, we assume that all potential curves can be described by the Morse function

\[
V(r_{\text{OH}}) = D_0 [1 - \exp(-a(r_{\text{OH}} - r_e))]^2
\]

where \( r_e \) is the equilibrium value of \( r_{\text{OH}} \). This potential model contains two parameters, which may be expressed in terms of the first and third derivatives at equilibrium, \( k_2 = V''(r_e)/2 \) and \( k_3 = V'''(r_e)/6 \), respectively. One easily shows that \( a = -k_3/k_2 \) and \( D_0 = (k_3^3/(k_2^3))^1/4 \). The energies \( E_n \) for the Morse oscillator are known exactly, and can be used to calculate the frequency \( \nu_{\text{OH}} = (E_1 - E_0)/\hbar \), where \( \hbar \) is the Planck constant, yielding

\[
\nu = \omega \left( 1 - 2 \frac{\hbar \omega}{D_0} \right).
\]

Here \( \omega \) is the harmonic frequency defined by \( 2\pi\omega = \sqrt{2k_2/m} \), as can also be seen by putting \( k_3 \) equal to zero \( (m \) is the reduced mass of the oscillator).

The total potential energy curve is then represented by the sum of an unperturbed gas-phase part augmented with a small perturbation. Since the perturbation can be represented by a quadratic polynomial, we may set \( k_{2\text{ext}}^\text{bound} = k_2^o \) and \( k_{2\text{ext}}^\text{bound} = k_2^o + \Delta k_2 \) with \( \Delta k_2 \) being small. Notice that \( k_2^o \) is half the second derivative of the gas-phase potential frequency shift, at the \( r_{\text{gas}} \) value, etc. To first order in the small parameter we then get that the anharmonic frequency shift \( \Delta \nu = \nu_{\text{bound}} - \nu^o \) is given by

\[
\Delta \nu \approx \left[ \nu^o + 4(\nu^o - \nu^o) \right] \cdot \frac{\Delta k_2}{2k_2^o}.
\]

An expression for \( \Delta k_2 \) can be obtained by applying a second-order perturbation to the \( V_{\text{gas}} \) potential energy curve, defined in eqn (1a) in the main text.

\[
\Delta k_2 = -\frac{3}{2k_2^o} \frac{k_{1\text{ext}}^o + k_{2\text{ext}}^o}{k_2^o}.
\]

The expression for the harmonic shift, \( \Delta \nu \) is obtained by putting the second term between brackets equal to zero, or, equivalently, by setting \( \nu^o \) equal to \( \nu^o \). Since \( k_2^o \) is positive for both \( \text{OH}_w \) and \( \text{OH}_w^- \) and \( k_3^o \) is negative, we conclude that both contributions to \( \Delta k_2 \) have the same sign. As long as \( k_{2\text{ext}}^o \) and \( k_{2\text{ext}}^o \) have the same sign, one may say that either of them determines the sign of \( \Delta k_2 \). With the first contribution to \( \Delta k_2 \) being the largest, we conclude that the sign of \( \Delta k_2 \) determines the sign of \( \Delta k_2 \). Moreover, since the factor between square brackets in the expression for \( \Delta \nu \) is always positive, we conclude that indeed the sign of \( k_{1\text{ext}}^o \) determines the sign of the gas-to-surface frequency shift.

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Conflicts of interest

There are no conflicts to declare.
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