Optical imaging of surface chemistry and dynamics in confinement

Carlos Macias-Romero, Igor Nahalka, Halil I. Okur, Sylvie Roke

We imaged the interfacial structure and dynamics of water in a microscopically confined geometry, in three dimensions and on millisecond time scales, with a structurally illuminated wide-field second harmonic microscope. The second harmonic images reported on the orientational order of interfacial water, induced by charge-dipole interactions between water molecules and surface charges. The images were converted into surface potential maps. Spatially resolved surface acid dissociation constant (pK_{\text{a,water}}) values were determined for the silica deprotonation reaction by following pH-induced chemical changes on the curved and confined surfaces of a glass microcapillary immersed in aqueous solutions. These values ranged from 2.3 to 10.7 along the wall of a single capillary because of surface heterogeneities. Water molecules that rotate along an oscillating external electric field were also imaged.

Microscopic and nanoscopic structural heterogeneities, confinement, and flow critically influence surface chemical processes in electrochemical, geological, and catalytic reactions (1–5). Recent advances in micro- and nanotechnology are driven by these phenomena, often in combination with electrostatic field gradients that can be surface-intrinsic or externally applied. Micro- and nanocapillaries in combination with interfacial voltage gradients are used in the fabrication, manipulation, and characterization of droplets, interfaces (6–9), and highly amorphous materials (10). An electrostatic field gradient in a narrow channel can be used to separate (11) and identify analytes (12). The increase in complexity of new materials and nano- and microtechnological developments (13) requires technology that can track, in real time, three-dimensional spatial changes in the molecular structure of confined systems, such as curved interfaces and pores, as well as the response to electrostatic fields of these systems.

We have constructed a high-throughput, structured-illumination second harmonic (SH) microscope to perform real-time three-dimensional (3D) chemical imaging of surfaces and confined geometries. Nonresonant coherent SH generation occurs only when noncentrosymmetric molecules, such as water, are oriented noncentrosymmetrically (14). This selection rule allows for probing just a few monolayers at an interface. We used this property to image the orientational order of water molecules at the inner and outer surfaces of a glass microcapillary immersed in aqueous solution. The surface chemistry of the microcapillary could be altered by changing the bulk pH of the solution. The images were converted into maps of the surface potential and surface chemical equilibrium constants. Although average values of the surface acid dissociation constant pK_{\text{a,water}} agree with previously reported values, structural interfacial heterogeneity was apparent, with pK_{\text{a,water}} values ranging locally from 2.3 to 10.7. Dynamical structural changes induced by an external oscillating electrostatic field were also imaged on millisecond time scales and in three dimensions. The observed structural fluctuations at the interface and in the tip opening (diameter <1 \mu m) originate from water molecules that interact with the external electrostatic field.

The wide-field (15–17) structured-illumination/HiLo (18) SH microscope is shown schematically in Fig. 1A. To achieve the wide-field geometry, we used a cosine diffraction phase grating generated with a spatial light modulator (SLM) to split the illuminating pulse beam (1036 nm, 200 kHz, 168 fs) into two beams. The SLM was then imaged on the focal (sample) plane of the objectives with the aid of a relay system. The field of view, 150 \mu m (full width at half maximum), was determined by the spot size on the SLM and the overall magnification of the relay system. The spatial frequency \( \sigma \) of the diffraction grating determined the angle of incidence of the beams on the sample. The image was recorded with an electron-multiplying intensified charge-coupled device (EM-ICCD) camera. Nonlinear polarimetry (19) was performed by controlling and analyzing the polarization state of the illuminating and emitted beams. A polarization state generator, comprising a half- and a quarter-wave plate, was used. The polarization state of the emitted light was analyzed with a half-wave plate placed in the emission path, followed by a polarization beam splitter (20).

In a wide-field illumination geometry, 3D imaging is not possible because the beams are too wide. To achieve 3D imaging and to improve the transverse spatial resolution (21), we used an adapted HiLo imaging procedure (18, 22–27) that relies on partial spatial coherence introduced by a chirp in the illuminating pulse. The procedure is based on Fourier filtering using a cosine intensity pattern containing a carrier spatial frequency to reject out-of-focus light (20). With this pattern, structured images \( I_n \) and \( I_c \) were produced, the latter being phase-shifted by \( \pi/2 \). The processed image was obtained with a uniformly illuminated image \( I_u = I_e + I_o \) and the structured-illumination image \( I_d \) using a HiLo algorithm (18, 23, 26–29).

We imaged a glass microcapillary immersed in an aqueous solution of pH-neutral ultrapure water with 10 mM NaCl. A phase-contrast image is shown in Fig. 1B; the structured SH images \( I_n \) and \( I_c \) are shown in Fig. 1, C and D. The uniformly illuminated image obtained from \( I_d = I_e + I_o \) is shown in Fig. 1E, and using the algorithm of Mertz and Kim (27), we obtained the final HiLo image of Fig. 1F (20), where background rejection leads to much better contrast. The corresponding spectrum consisted of a single peak centered at 515 nm (Fig. 1A, left inset), confirming that the image contrast originates from SH emission. The SH intensity was detected in the XXX polarization combinations (i.e., with all beams polarized and analyzed in a direction perpendicular to the main symmetry axis of the microcapillary) (Fig. 1C). Light polarized along the interface did not generate SH photons. The measured transverse resolution was 188 nm (Fig. 1A, half width at maximum (HWWM) of the red curve in the right inset), as measured by imaging BaTiO\(_3\) particles 50 nm in diameter (30) on a glass coverslip. The best possible axial resolution was 520 nm for two-photon fluorescence (fig. S2) (20). The imaging throughput was improved by a factor of \( \sim 5 \times 10^3 \) relative to a scanning confocal two-photon microscope (31) and to our previous 2D wide-field SH microscope (20, 32).

Surface chemical changes were SH-imaged and induced on three different glass microcapillaries in solution with pH values of 2, 7, and 12. The ionic strength was kept constant by adding 10 mM NaCl. The silica surface in contact with a solution of pH = 2 is on average charge-neutral, with the vast majority of the silanol groups in the protonated SiOH state (33, 34). Increasing the bulk pH of the solution increased the fraction of surface SiOH groups, resulting in a predominantly negatively charged surface with average charge densities of \( \sim 2 \mu \text{C/cm}^2 \) under pH-neutral conditions and \( \sim 17 \mu \text{C/cm}^2 \) at pH = 12 (33, 36). Increasing the net surface charge caused the interfacial electrostatic field \( E_{\text{DC}} \) to grow in magnitude, and the interaction of this field with the dipole moment of water distorted the orientational distribution of water molecules. The dipole moment of the water molecules aligned with \( E_{\text{DC}} \), which allowed SH photons to be generated for probing aqueous silica interfaces (35, 36). The spatially averaged and spatially averaged spectroscopic SH measurements have shown that the breaking of centrosymmetry only a few nanometers from the interface is sufficient for generating a detectable SH intensity. The spatially and temporally integrated SH intensity can be correlated to the surface potential \( \Phi_0 \) according to \( I(2\omega) \sim \left| \chi_{\text{xx}}^{(2)} + \chi_{\text{yy}}^{(3)} \Phi_0 f_0 \right|^2 \) (35, 37), which originates from

\[
I(2\omega; x, y, t) = I(\omega; x, y; t) \left| \chi_{\text{xx}}^{(2)}(x, y, t) + \chi_{\text{yy}}^{(3)} f_0 E_{\text{DC}}(x, y, z) \right|^2
\]

\[t \geq \tau_i\]

\[
\chi_{\text{yy}}^{(3)} f_0 \int_0^\infty E_{\text{DC}}(x, y, z) dz\]

\[t \geq \tau_i\]

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where \( x, y, z \) represents the local surface coordinate system (with \( z \) being the surface normal), \( E_{DC}(x, y, z) \) relates to the electrostatic potential \( \Phi \) via \( -\nabla \Phi(x, y, t) \), \( \chi^{(2)} \) is the second-order susceptibility of the capillary silica/water surface, and \( f_2 \) is an interference term that takes the value 1 for SH transmission experiments (37). \( \chi^{(3)} \) is an effective third-order susceptibility that contains different contributions (20); the largest is the reorientation of water molecules in the electric double layer ([37], here ~99%) (20). The response in Eq. 1 can be used to determine changes in the surface potential \( \Phi_0 \).

The average SH intensity increased in the order pH(2) < pH(neutral) < pH(12), in agreement with Macias-Romero et al., Science 357, 784–788 (2017).
Fig. 3. Dynamic imaging of surface chemical changes. (A) Illustration of the experiment.
(B) Illustration of the applied field lines (ΔV > 0). (C) Computed ionic charge density in the aqueous solution adjacent to the interface (fig. S7) (20). (D) Electrostatic field and potential drop for ΔV = 10 V and ΔV = −10 V along the symmetry (Y) axis of the capillary. (E) Snapshots of movie S1, recorded with acquisition time of 250 ms for different values of the applied potential (i, 10 V; ii, −10 V; i + ii, composite of five frames each). The ΔV value corresponding to i, ii, iii, and iv is shown in (G). The legend displays the color coding associated with the applied potential, polarization combination, and multiplication factor. The inset at lower right displays the tip region only. All images were normalized with respect to the illuminating beam profile. (F) 3D rendering of the 10 V and −10 V channels (45 stacks per 4° rotation around the Y axis). The inset shows a zoom-in of the tip. (G) ΔV (green), the spatially integrated SH intensity (XXX polarization, red), and the numerical computation (blue) (20).

expectations from spectroscopic measurements (Fig. 2A) (35, 38–41). We converted the SH intensity in the images of Fig. 2A to a change in the surface potential \( \Phi_0 \). Although it was not possible to determine absolute values for the surface potential in this experiment (37), assuming that \( \Phi_0 = 0 \) at pH = 2 and that the average charge density under pH-neutral conditions is −2 µC/cm² (33, 34), we constructed a scale for \( \Phi_0 \) using the Gouy-Chapman-Stern model to provide reference values (36). The retrieved values are given by the color bar in Fig. 2A. The average values of \( \Phi_0 \) agree well with reported values (34, 38–40). However, the variation in the spatial distribution of intensities/surface potentials was significant, reaching values of ~3–4 \( \Phi_0 \). This variation reflected the diverse values reported (34, 38–40) and indicated that the surface was structurally not uniform—for example, in terms of the distribution of charged SiO⁻ or the nanoscopic surface geometry imposed by the production procedure of the capillary (20). Indeed, mono-, bi-, or trimodal deprotonation behavior with surface pKₐ values of 6.8 (42), 4.8, 8.5 (35), or 3.8, 5.2, and ~9 as well as a dependence on the pH history of the sample (40) have been reported. Computations show that the local chemical structure and hydrogen-bonding environment, as well as strains or defects in the surface structures, all contribute to heterogeneity (43, 44).

We determined the spatial distribution of pKₐ values for the =SiOH⁻ + H₂O \( \rightarrow \) SiO⁻ + H₃O⁺ reaction by flowing a pH = 12 solution through a capillary that was charge-neutral on average (Fig. 2, B to D), using a shear rate of 8.5 × 10⁻² s⁻¹. We imaged surface potential changes every 250 ms, while the bulk pH changed in the direction of the arrow in Fig. 2C. The pH of the solution was determined with a pH indicator dye solution (20) in a separate experiment. Knowing the bulk pH value inside the capillary, we determined the pKₐ value of every pixel according to

\[
K_{a,s} = \frac{H_2O_{(b)}}{H_2O_{(s)}} \exp \left( \frac{-\Phi_0}{k_B T} \right),
\]

and

\[
pK_{a,s} = -\log(K_{a,s}).
\]

where \( N_{-SiOH} / N_{-SiO} \) is the ratio of deprotonated over protonated groups, which can be derived from \( \Phi_0 \) using Gouy-Chapman-Stern theory (20, 45).

Figure 2B shows the distribution of pKₐ values for the deprotonation reaction, integrated along the X direction. Figure 2C displays the surface potential and derived corresponding surface charge density for two pixels as a function of pH (with pKₐ(A) = 3.8 and pKₐ(B) = 5.9). Figure 2D displays histograms of the obtained pKₐ values with an average of 6.7. The chemical reactivity varied spatially from 2.3 < pKₐ < 10.7 and followed a Gaussian distribution. The average value agreed with that in (42), but substantial variations in chemical reactivity across the surface were observed, which would explain why different studies (even using the same methods (35, 40)) have found different average pKₐ values.

To image additional types of dynamical changes at interfaces and in confinement, we placed two Ag/AgCl electrodes in a pH-neutral solution that also contained the microcapillary and 10 mM NaCl (Fig. 3A). The electrode outside the capillary was grounded, and the applied electrostatic potential difference was varied sinusoidally on the inner electrode from 10 V to −10 V at 0.05 Hz (−1 V/s). The spacing between the electrodes was 1 cm, and we imaged the 60-µm-long edge, which had an opening with diameter <1 µm. Solving numerically the Navier-Stokes, Nernst-Planck, and Poisson equations by means of a COMSOL routine, we computed the resulting average changes in E_{max} as well as the induced flow in the aqueous solution (20, 46) (Fig. 3B). The electrostatic field strength was high (>10⁸ V/m) (fig. S9) in the interfacial region and in the tip of the microcapillary. Figure 3C displays the computed average ionic charge density close to the surface for different values of ΔV (using an initial surface charge density of −2 µC/cm² (20, 33, 34)). As ΔV was cycled from 10 V to −10 V, the electrostatic field at the surface (z = 0) of the microcapillary changed as the distribution of Na⁺ and Cl⁻ ions in the electric double layer changed (fig. S9).

Water molecules that are dynamically oriented by the interfacial DC field can be probed with the XXX polarization combination, as in Fig. 2. Another place where the electrostatic field reaches similarly high values is inside the opening of the microcapillary (diameter <1 µm) (46). As plotted in Fig. 3D, here E_{max} oscillates between 0.75 × 10⁶ and −1.15 × 10⁶ V/m, which corresponds to a fraction of 5 × 10⁻¹ oriented water molecules that are fully aligned by it (fig. S4A), or, equivalently, ∼1 million aligned water molecules per pixel. These aligned water molecules were probed with the YYY polarization combination (i.e., with all beams polarized and analyzed along the main symmetry axis of the capillary).

To image the change in the orientational order of water molecules in both regions as a function of time and applied potential, we recorded 3D SH images (integration time, 250 ms) in different polarization combinations (XXX to probe the interfacial water and YYY to probe the water in the tip). Movie S1 shows the temporal evolution of the SH intensity. Figure 3E shows 250-ms snapshot cross sections for ΔV = 10 V (i), −10 V (ii), and their composite (averaging 5 frames). The acquisition time, 250 ms, was sufficient to resolve the interfacial water as well as the −10⁶
oriented water molecules in the tip. Figure 3F shows a 3D rendering constructed from 45 image stacks, with each image taken at 4° increments. Movie S2 shows a composite 3D rendering of the oriented water at 10 V (red, cyan) and −10 V (green, purple). There was relatively more oriented water on the inside surface for ΔV > 0 (Fig. 3E, movie S2, and fig. S11), which agrees with the higher Epc there (Fig. 3C and fig. S9). Figure 3G shows the applied potential difference ΔV′ (green), the spatially averaged SH intensity of the interfacial water (red, XXX polarization), and a numerical computation of the intensity using Eq. 1 (blue) (20). Good agreement was achieved between the mean-field model and the averaged SH intensity changes. However, the surface response in the images of Fig. 3E (or movie S1) shows that the intensity distribution in the images was not uniform. This difference points toward local variations in the surface structure, potential, and chemical reactivity similar to those observed in Fig. 2 (20).

We imaged dynamical interfacial changes as well as small amounts of oriented confined water in three dimensions and on millisecond time scales. The real-time observation of heterogeneous or transient structures should prove valuable for the development of micro- and nanotechnology that critically relies on local structural and dynamical processes (I1), but also for geochemistry, catalysis, electrochemistry such as in fuel cells (5, 11, 13, 47), and processes using electrokinetic phenomena in micro- and nanofluidic environments (48). The imaging of the orientational order of water in confinement could be used to noninvasively infer the structure of macromolecules, such as lipids, DNA, and proteins, or structural changes induced by potential gradients.

REFERENCES AND NOTES

20. See supplementary materials.
22. Note that HIL0 microscopy is different from HIL0 (highly inclined and laminated optical) sectioning microscopy.
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Imaging surfaces with water
The surfaces of real materials are often highly chemically heterogeneous, and the reported values of even simple properties such as surface acidity can vary widely in many cases. Macias-Romero et al. developed a microscope that images surfaces on the basis of second-harmonic generation from the orientation of interfacial water (see the Perspective by Hunger and Parekh). They followed the deprotonation of silica along glass micropipettes by changing solution pH and found many regions where the surface acidity deviated strongly from the average for the entire micropipette.

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