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Electromechanical imaging of biological systems with sub-10 nm resolution

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Electromechanical imaging of tooth dentin and enamel has been performed with sub-10 nm resolution using piezoresponse force microscopy. Characteristic piezoelectric domain size and local protein fiber ordering in dentin have been determined. The shape of a single protein fibril in enamel is visualized in real space and local hysteresis loops are measured. Because of the ubiquitous presence of piezoelectricity in biological systems, this approach is expected to find broad application in high-resolution studies of a wide range of biomaterials. © 2005 American Institute of Physics.

Here, we demonstrate electromechanical imaging and microstructural analysis of dentin and enamel in a human tooth with sub-10 nm resolution using vertical and lateral piezoresponce force microscopy (PFM). In PFM, a conductive tip is brought into contact with the surface, and the piezoelectric response is detected as the first harmonic component. $A_{1\omega}$ of the tip deflection, $A=A_0+A_1\omega \cos(\omega t+\varphi)$, induced by the application of the periodic bias $V_{tip}=V_{dc}+V_{ac}\cos(\omega t)$ to the tip. Principles and image formation mechanism of PFM are described in detail elsewhere. PFM is implemented on a commercial scanning probe microscopy (SPM) system (Veeco MultiMode NS-III) equipped with additional function generators and lock-in amplifiers (DS 345 and SRS 830, Stanford Research Instruments, and Model 7280, Signal Recovery). A custom-built sample holder was used to allow direct tip biasing and to avoid capacitive cross talk in the SPM electronics. Measurements were performed using Pt- and Au-coated tips (NCSC-12 C, Micromasch, $l=130\mu m$, resonant frequency $\sim150$ kHz, spring constant $k=4.5$ N/m). Vertical PFM (VPFM) measurements were performed at frequencies 50–100 kHz, which minimizes the longitudinal contribution to measured vertical signal. For lateral PFM (LPFM), the optimal conditions for contrast transfer were $\sim10$ kHz; for higher frequencies, the onset of sliding friction minimizes in-plane oscillation transfer between the tip and the surface. Custom LABVIEW software was developed for simultaneous acquisition of VPFM and LPFM phase and amplitude data, emulating additional SPM data acquisitions channels.

The surface layer of tooth enamel is comprised primarily of hydroxyapatite (HAP) crystals and a small fraction ($\sim3\text{–}5\%$) of organic fibers concentrated mostly in the vicinity of the dentin-enamel junction (DEJ). The dentin layer below the enamel has significantly higher fraction of organic...
material, up to 30–40%. The dominant organic constituent of dentin and other calcified and connective tissues is collagen. Collagen, the most abundant protein in biological systems, exhibits piezoelectric properties, resulting in the nearly ubiquitous presence of piezoelectricity in connective and partially calcified tissues.13

A deciduous human tooth was cross sectioned parallel to growth direction and polished using diamond polishing pads down to 0.5 μm grit size. The enamel and dentin regions can be readily identified using optical microscopy. Surface topography of dentin and enamel regions is shown in Figs. 1(a) and 1(b).14 It can be seen that both regions are comprised of elongated grains of about 100–300 nm in size, however, the difference in microstructure between the two regions is small. Shown in comparison are simultaneously acquired amplitude and phase VPFM images [Figs. 1(c)–1(f)]. In enamel, several isolated regions of 50–200 nm in size with a high piezoresponse signal (appearing as bright spots in amplitude image) are observed, while the majority of material is nonpiezoelectric. At the same time, most of the dentin region exhibits a strong response signal, with the typical size of regions of constant phase and amplitude (piezoelectric domains) of the order of 200 nm. These observations are consistent with a high density of piezoelectrically active proteins, such as collagen, in dentin13 and a low fraction of proteins in enamel.

Simultaneous measurements of the vertical and lateral piezoresponse map allow us to determine two components of local electromechanical response vector, which can be further correlated with topography or elasticity data from, e.g., phase atomic force microscopy. Particularly, VPFM and LPFM data provide the basis for the statistical description of the electromechanical microstructure of the material. Shown in Fig. 2(a) is the double histogram of normalized VPFM and LPFM signals obtained in the dentin region, representing the count number of points with the signal level in the interval \( \langle \text{vpr} + \Delta v, \text{lpr} + \Delta l \rangle \), where \( \text{vpr}, \text{lpr} \in (-1, 1) \). Shown in Figs. 2(b) and 2(c) are the amplitude, \( A_{2D} \), and angle, \( \theta_{2D} \), signal distributions, where \( A_{2D} = |\text{vpr} + \text{lpr}| \), \( \theta_{2D} = \text{Arg}(\text{vpr} + \text{lpr}) \) calculated using commercial image analysis software.15 Note that the angle calculated from VPFM and LPFM data represents the orientation of the piezoresponse vector in the plane perpendicular to the cantilever axis and thus serves as the measure of local collagen fibril orientation.

Data shown in Figs. 2(a) and 2(c) illustrate that there are two primary antiparallel orientations of the piezoresponse vector. Thus, the local dentin microstructure can be well represented by axially ordered antiparallel collagen fibers, as shown in the inset of Fig. 2(d). The characteristic fiber size can be determined using self-correlation function analysis as is illustrated in Fig. 2(d). The normalized experimental function can be well approximated using a simple phenomenological form \( C(x) = A \exp(-x/\xi) \), where characteristic domain size \( \xi \) is 160±2 nm, in agreement with data from Fig. 1(d).

To get further insight into the structure of the isolated protein (presumably amelogenin) fibrils in enamel, we have performed high-resolution VPFM and LPFM imaging of the enamel region in the DEJ vicinity, as illustrated in Fig. 3. Both VPFM and LPFM images show a very strong electromechanical response that we attribute to a protein fibril embedded within a nonpiezoelectric matrix. Comparison of the VPFM and LPFM images shows different patterns of piezoelectric domains, suggesting a complicated structure of the protein fibril, consisting of several protein molecules. Notably, the spatial resolution of PFM, determined as a half-width of the boundary between differently oriented piezoelectric regions, is ~5 nm. This is comparable to the best results achieved to date for thin films of ferroelectric perovskites and well above ~30–100 nm resolution for single crystals.
Finally, to check the presence of the ferroelectric polarization in the material, PFM was used to acquire local hysteresis loop in a manner similar to ferroelectric materials.\footnote{E. Fukada and I. Yasuda, J. Phys. Soc. Jpn. \textbf{12}, 1158 (1957).} The piezoresponse, measured as a function of a dc bias offset on the tip, is shown in Fig. 4. No inversion of the strain sign upon application of the dc bias was observed, indicating that collagen molecules possess strong piezoelectric properties, comparable to quartz, but are not ferroelectric. Notably, the slope of the amplitude response is small, indicative of minimal electrostatic contribution to measured signal. The effective piezoelectric coefficient is $d_{\text{local}} = 0.15\text{--}0.25 \ \text{pm/V}$, nearly ten times larger than $d = 0.028 \ \text{pm/V}$ for a macroscopic dentin sample and comparable to $d = 0.28 \ \text{pm/V}$ for dry bone.\footnote{A. A. Marino and B. D. Gross, Arch. Oral Biol. \textbf{34}, 507 (1989).} Interestingly, these observations are consistent with the antiparallel orientation of collagen fibrils in dentin, since the electromechanical response will be cancelled out on the macroscopic scale.

To summarize, we have demonstrated the electromechanical imaging of dental tissues with sub-10 nanometer resolution. PFM provides an approach to distinguish between materials with different electromechanical properties, such as proteins and calcified tissues. PFM allows statistical description of dental tissue structure, including the characteristic fiber size and type of local ordering. The collagen fibril has been visualized in real space. The local electromechanical properties of dental tissues are measured as a function of tip bias and the material is shown to be piezoelectric, rather than ferroelectric. Due to the ubiquity of piezoelectricity in biological systems, we expect this approach to find broad application in future high-resolution studies of biomaterials.

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\begin{figure}
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\includegraphics[width=\textwidth]{fig4}
\caption{Piezoelectric hysteresis loops of a single-collagen fibril in tooth enamel.}
\end{figure}