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# Microwave Microscopy and Its Applications

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microwave microscopy, 2D semiconductors, quantum materials, topological edge states, domain walls, microwave acoustics

## Abstract

Understanding the nanoscale electrodynamic properties of a material at microwave frequencies is of great interest for materials science, condensed matter physics, device engineering, and biology. With specialized probes, sensitive detection electronics, and improved scanning platforms, microwave microscopy has become an important tool for cutting-edge materials research in the past decade. In this article, we review the basic components and data interpretation of microwave imaging and its broad range of applications. In addition to the general-purpose mapping of permittivity and conductivity, microwave microscopy is now exploited to perform quantitative measurements on semiconductor devices, photosensitive materials, ferroelectric domains and domain walls, and acoustic-wave systems. Implementation of the technique in low-temperature and high-magnetic-field chambers has also led to major discoveries in quantum materials with strong correlation and topological order. We conclude the review with an outlook of the ultimate resolution, operation frequency, and future industrial and academic applications of near-field microwave microscopy.

## 1. INTRODUCTION

The nanoscale electrodynamic response of solid-state materials is of fundamental importance for research on novel electronic devices with nanometer-sized features, ferroic crystals with mesoscopic domains and domain walls (DWs), strongly correlated materials with spatially inhomogeneous phases, and topological systems with distinct edge and bulk properties. In the far-field regime, however, the result of a measurement is always averaged over a length scale defined by the electromagnetic (EM) wavelength (1). Going beyond this diffraction limit would demand the use of a sharp near-field probe in close proximity with the specimen, usually implemented on a scanning probe microscopy (SPM) platform. In the infrared and visible range of the EM spectrum, such near-field scanning optical microscopy (NSOM) has greatly expanded our knowledge on the mesoscale optical properties of functional materials (2–4). In recent years, the analog of NSOM in the microwave regime, or near-field scanning microwave microscopy (NSMM), has also gained tremendous momentum (4–7). In most cases, the interaction between microwave radiation and materials is much more classical than the optical counterpart. The relative simplicity in the design of NSMM and straightforward data interpretation have led to new types of spatially resolved experiments that are otherwise not feasible.

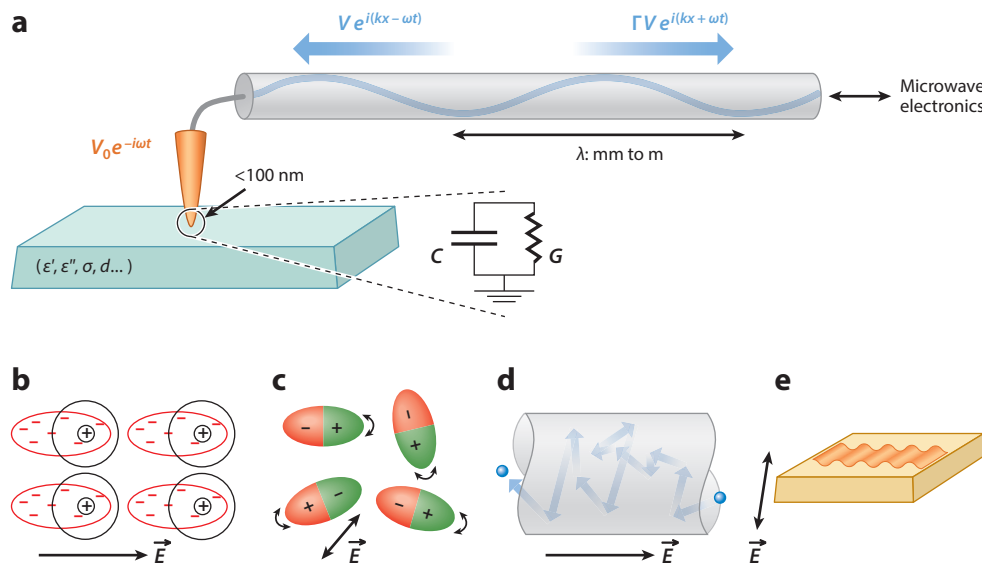
NSMM was first envisioned in 1928 (8), followed by a few early prototypes (9–12). With the invention of modern SPM, most notably scanning tunneling microscopy (STM) (13) and atomic force microscopy (AFM) (14), various designs of NSMM were demonstrated in the 1990s (15–20), and the focus was mainly on the understanding of the contrast mechanism, the modeling of tip-sample interaction, and the system calibration using standard dielectrics and semiconductors. Excellent reviews of these efforts can be found in References 4–6. In the past decade, much progress was made to integrate microwave probes with state-of-the-art scanning platforms (21–26) and low-temperature/high-magnetic-field chambers (27). Propelled by the rise of quantum materials that are inherently heterogeneous, NSMM has evolved into an important SPM mode for cutting-edge scientific research. While the technique continues to develop and new applications continue to emerge, it is now appropriate to provide a work-in-progress review on the general principle of microwave microscopy, its deployment in materials research, and future prospects for the years to come. We would like to point out that this article is by no means a comprehensive summary of the vast literature in this field, for which the readers are referred to other reviews (4–6). It is rather a survey of the key works and a call for collective efforts to advance this burgeoning area of research.

The rest of this review is organized as follows. In Section 2, we begin by reviewing the basics of microwave microscopy, followed by a description of the system design and data analysis. We then elaborate its applications, categorized by the different material properties probed by the technique, in Section 3. For each subsection, we also discuss the specific targets in the near term. We conclude the review in Section 4 with a summary of the current status and an outlook of this area for the next few decades.

## 2. NEAR-FIELD MICROWAVE MICROSCOPY

### 2.1. Basic Principles

Microwaves are commonly defined as the EM radiation with free-space wavelength  $\lambda$  ranging from 1 m (frequency  $f = 300$  MHz) to 1 mm ( $f = 300$  GHz), although we focus on a narrower band from 300 MHz to 20 GHz here. The interaction between matter and the EM wave in this regime is well understood and widely exploited in day-to-day technology—to name a few, food



**Figure 1**

Schematics of microwave impedance microscopy and its contrast mechanism. (a) Conceptual sketch of the microwave microscope, showing the excitation and reflection waves in the transmission line, the quasi-static voltage at the nanoscale tip, and the material properties to be probed. The lumped elements representing the tip-sample interaction are shown in the inset. (b–e) Illustrations of (b) the dielectric response of an insulator to an  $E$ -field, (c) dipolar oscillation in the microwave field, (d) diffusive conduction of an electron, and (e) excitation of acoustic waves in a piezoelectric crystal by an alternating  $E$ -field.

is heated in a microwave oven while a microwaveable plate remains lukewarm; an airplane made of metal, on the other hand, reflects microwaves and appears on the radar screen. While sharing the same characteristics, NSMM differs from these phenomena in various aspects. First, while the electric and magnetic parts of the EM energy density are comparable in propagating waves, only one is dominant in each NSMM design (6). In **Figure 1a**, it is the gigahertz electric field ( $E$ -field) that is maximized at the electrically open tip. Furthermore, as the typical tip size is approximately six orders of magnitude smaller than the wavelength, the tip-sample interaction can be accurately described by the lumped elements in **Figure 1a**. Such electrical-impedance-type NSMM is referred to hereafter as microwave impedance microscopy (MIM) (7, 28). Finally, the quasi-static fields only extend into the sample with a dimension determined by the tip apex. To form a near-field image, therefore, one must raster scan the tip on the sample surface.

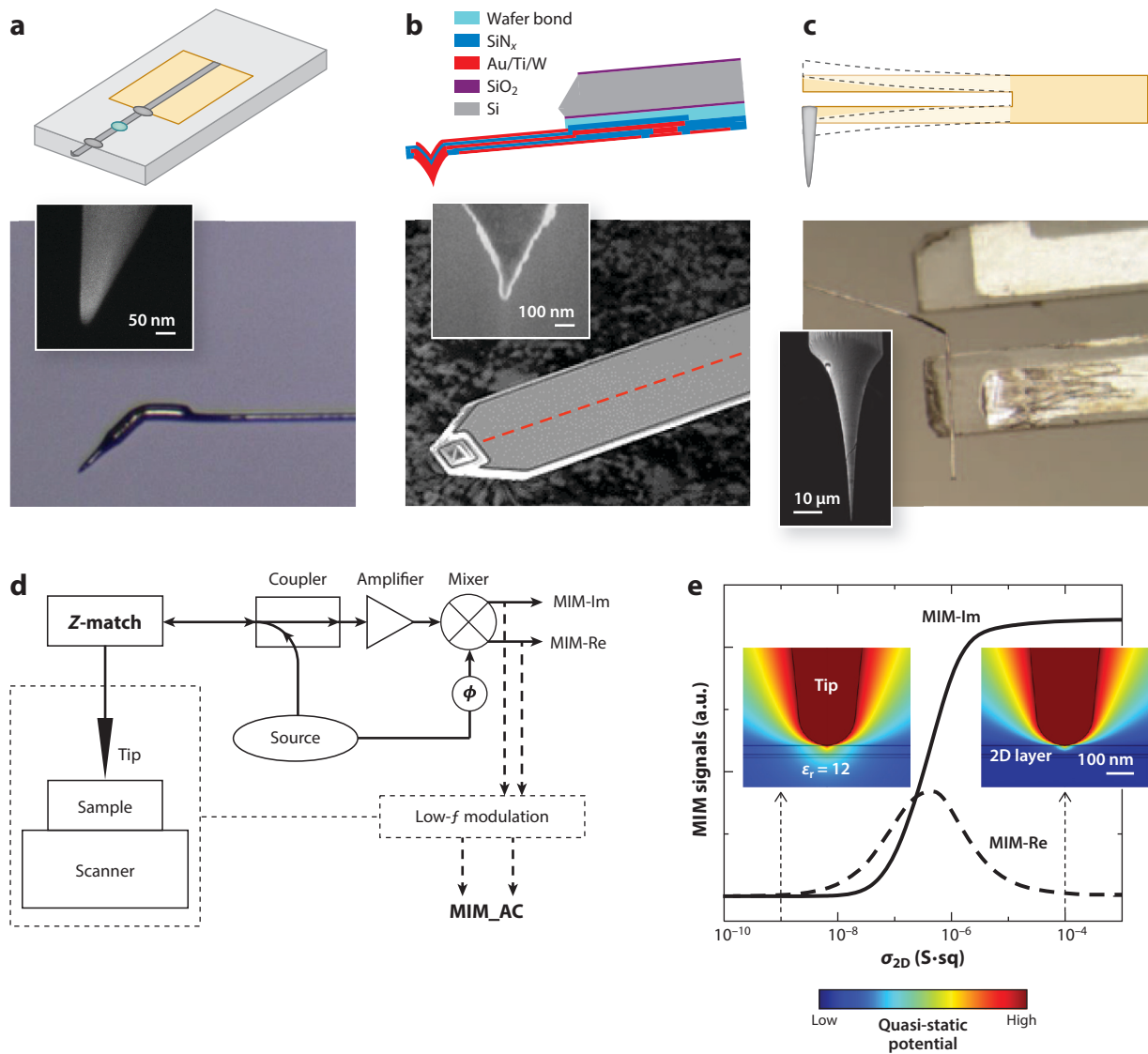
Illustrations of various MIM contrast mechanisms are shown in **Figure 1b–e**. The electrical response of a material can be expressed as  $\epsilon' + i(\epsilon'' + \sigma/\omega)$ , where  $\epsilon'$  and  $\epsilon''$  are real and imaginary parts of the permittivity,  $\sigma$  is the conductivity, and  $\omega = 2\pi f$  is the angular frequency. As depicted in **Figure 1b**, the response of a perfect insulator to gigahertz  $E$ -fields is purely capacitive since the displacement current is  $90^\circ$  out of phase with respect to the tip voltage. For materials with permanent dipoles or gigahertz collective modes, the interaction with an oscillating  $E$ -field leads to dielectric loss (**Figure 1c**). For electrical conductors, the motion of conduction electrons (**Figure 1d**) will also result in energy dissipation. Finally, in piezoelectric samples, the electromechanical transduction into gigahertz acoustic waves leads to another channel of energy loss (**Figure 1e**). Other less common configurations of microwave microscopy are briefly discussed in Section 3.5.

## 2.2. System Components

The schematic in **Figure 1a** is, of course, oversimplified for MIM. The actual system would at least consist of the probe, detection electronics, and the scanning platform. Although metalized tips with a nanoscale apex are readily available for conductive atomic force microscopy (C-AFM), they are unsuitable for MIM due to the strong stray-field contribution from the cantilever body. One design to mitigate this problem, as shown in **Figure 2a**, is to increase the sample-cantilever distance by a tall ( $\sim 80\text{ }\mu\text{m}$ ) shank of the metal tip (24, 25), which is commercially available from Rocky Mountain Nanotechnology, LLC. Here, the nanometer-sized apex is achieved by the chemical etching of Pt wires. A more elegant solution is the batch fabrication of shielded cantilever probes (29), which are commercially available from PrimeNano Inc. As indicated in **Figure 2b**, the Au/Ti/W center conductor is sandwiched between  $\text{SiN}_x$  layers and shielded by metal layers. The sub-100-nm tip apex is obtained by oxidation sharpening of the sacrificial Si pit and the subsequent metal refill. A drawback here is the inevitable wear during the contact-mode scans, which strongly affects the MIM signal level. In this regard, quartz tuning-fork (TF) probes with etched metal tips (**Figure 2c**) are desirable not only for the preservation of the tip apex but also for their self-sensing capability in a cryogenic environment (30, 31). The lack of shielding is again circumvented by the high-aspect-ratio probe. The distance modulation in a TF-MIM also rejects the background drift and provides absolute sample information (31). The disadvantages of TF probes are the much harder operation compared with cantilevers and the difficulty tracking rough surfaces due to the small oscillation amplitude (usually  $<10\text{ nm}$ ). Both types of probes will continue to be used in MIM research.

**Figure 2d** shows a schematic of a complete MIM system. Since the impedance of an electrically open tip is very different from  $50\text{ }\Omega$  in typical transmission lines, an impedance-match (Z-match) section is required to ensure efficient power transfer to the probe. Examples of Z-match include cavity resonators for needle probes (6) and  $\lambda/2$  or  $\lambda/4$  transmission-line resonators for cantilever (28) and TF (30) probes. In **Figure 2d**, the excitation signal is sent to the tip through a directional coupler and the reflected signal is amplified and demodulated by an in-phase-quadrature mixer, similar to the  $S_{11}$  measurement in a vector network analyzer (24, 25). The input microwave power is typically  $1 \sim 10^\circ\text{W}$ , and the measurement is mostly noninvasive (7, 28). As long as the load impedance is a small perturbation to the tip impedance, the imaginary microwave impedance microscopy (MIM-Im) and real microwave impedance microscopy (MIM-Re) parts of the output signals are proportional to the effective tip-sample capacitance and conductance (**Figure 1a**), respectively. A rigorous derivation of the MIM signals can be found in Reference 32. Finally, it is sometimes necessary to modulate the MIM output by a low- $f$  (usually kilohertz) signal, such as voltage modulation for semiconductor samples (33) and distance modulation in the TF operation (30, 31). In this case, lock-in amplifiers are needed to produce the MIM\_AC outputs, as depicted in **Figure 2d**.

A brief introduction to MIM data interpretation is provided at the end of this section. While analytical solutions of certain sample geometries can be obtained (34–37), it is now practical to directly compute the quasi-static admittance (inverse impedance) of an arbitrary tip-sample configuration using the finite-element analysis (FEA) method (7, 28). **Figure 2e** plots the simulated MIM signals as a function of the sheet conductance  $\sigma_{2D}$  of a two-dimensional (2D) layer buried 30 nm below the surface of a dielectric sample with  $\epsilon_r = 12$ . As a generic feature, the MIM-Im signal increases monotonically with respect to  $\sigma_{2D}$ , whereas the MIM-Re signal peaks at an intermediate  $\sigma_{2D}$  that coincides with the inflection point in MIM-Im. It is obvious that there is an optimal window of  $\sigma_{2D}$  determined by the operation frequency and sample parameters. For materials with large conductivity contrast, MIM-Im is advantageous because of the monotonic response.



**Figure 2**

Components of MIM and data interpretation. (a, top) Schematic of the solid metal probe with a contact pad. (a, bottom) The cantilever probe with a tall shank of the metal tip. The inset shows the apex of the etched Pt tip. (b, top) Layer structure of the shielded cantilever probe. (Bottom) SEM of the cantilever and pyramidal tip. The red dashed line indicates the buried center conductor. The inset shows the apex of the Au/Ti/W tip. (c, top) A tuning-fork-based probe. (Bottom) An etched W wire (SEM in the inset) glued to a tuning fork. (d) Schematics of the MIM setup. (e) Simulated MIM signals as a function of the sheet conductance of a subsurface 2D layer  $\sigma_{2D}$ . The insets on the left and right show the quasi-static potential distribution maps for  $\sigma_{2D} = 10^{-9}$  S·sq and  $10^{-4}$  S·sq, respectively. Abbreviations: MIM, microwave impedance microscopy; MIM-Im, microwave impedance microscopy–imaginary part; MIM-Re, microwave impedance microscopy–real part; MIM\_AC, microwave impedance microscopy–alternating current; SEM, scanning electron micrograph. Panel b adapted with permission from Reference 29; copyright 2012 IOP Publishing. Panel c adapted with permission from Reference 30; copyright 2016 AIP Publishing.

Near the insulating or conductive limits, however, MIM-Re is usually chosen due to the larger  $d(\text{MIM-Re})/d\sigma$  here and the small topographic cross-talk. This simple exercise also demonstrates that one cannot directly associate the real and imaginary parts of the dielectric function with the MIM-Re and MIM-Im signals, respectively. Finally, the quasi-static potential at the two limits of  $\sigma_{2D}$  is shown in the inset of **Figure 2d**. The spatial extent of the  $E$ -field (gradient of potential), which defines the spatial resolution of MIM, is comparable to the tip diameter.

### 3. APPLICATIONS OF MICROWAVE MICROSCOPY

NSMM has many applications in condensed matter physics, materials science and engineering, electrical engineering, and biology, as well as other interdisciplinary research. In the following sections, we group the applications in terms of the contrast mechanism enumerated in **Figure 1**, followed by brief discussions under each category.

#### 3.1. Imaging Dielectric Constant

In an electrical insulator, the ratio between the external  $E$ -field and the net field in the medium is defined as its dielectric constant or relative permittivity  $\epsilon_r$ . For advanced nanomaterials, the conventional measurement of  $\epsilon_r$  involves complicated device fabrication into the parallel-plate geometry. On the other hand, MIM provides a noninvasive method to characterize this fundamental property in the nanoscale.

**Figure 3a** shows the AFM and MIM ( $f = 1$  GHz) images of the layered semiconducting chalcogenide  $\text{In}_2\text{Se}_3$  (38), an important system for phase-change memory, thermoelectric, and photoelectric applications (39). The MIM-Im image exhibits distinct contrast over the mica substrate as a function of the flake thickness, i.e., the signal is lower than mica for a layer number ( $n$ ) of 3 and higher for  $n > 4$ . FEA simulation of the MIM data in **Figure 3b** shows that the effective  $\epsilon_r$  increases monotonically from two to six layers and saturates to the bulk value after  $n = 8$ , consistent with first-principles calculations (38). The  $n$ -dependent  $\epsilon_r$  is ubiquitous for van der Waals materials (40), and the methodology is valuable for this vibrant research field.

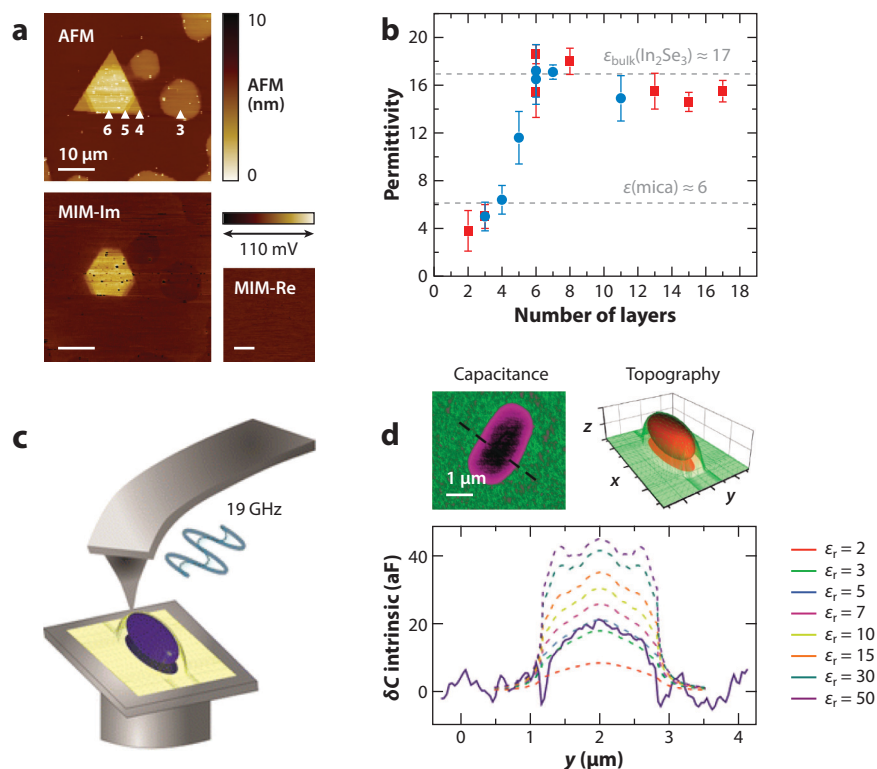
The gigahertz permittivity of biological systems is also of interest for medical applications as it determines the transmission, reflection, and absorption of microwaves in tissues (41). **Figure 3c** shows a schematic of microwave imaging on a single bacterial cell at 19 GHz (42). For the irregular cell structure, the morphologic contribution to the MIM-Im data cannot be ignored. As a result, it is crucial to perform careful calibration to remove the topographic cross-talk from the raw data (**Figure 3d**). The intrinsic capacitance after subtracting the background is plotted in **Figure 3d**, showing  $\epsilon_r \sim 4$  with no significant dielectric loss (42). The findings open an important avenue in label-free imaging of single cells with nanoscale spatial resolution.

The two examples reviewed here highlight the potential of MIM as a versatile permittivity imaging tool. Using standard dielectrics as calibration samples (24, 31, 43, 44), this method can be further applied to quantify  $\epsilon_r$  of nanoparticles, free-standing films, and granular ceramics, whose nanoscale dielectric response is otherwise difficult to probe.

#### 3.2. Imaging Electrical Conductivity

The imaging of local conductivity is arguably the most successful application of microwave microscopy. In particular, the ability to visualize subsurface conduction is of great interest across many research disciplines. In this section, we first introduce several examples of general-purpose conductivity imaging and then proceed with studies in more specialized areas.

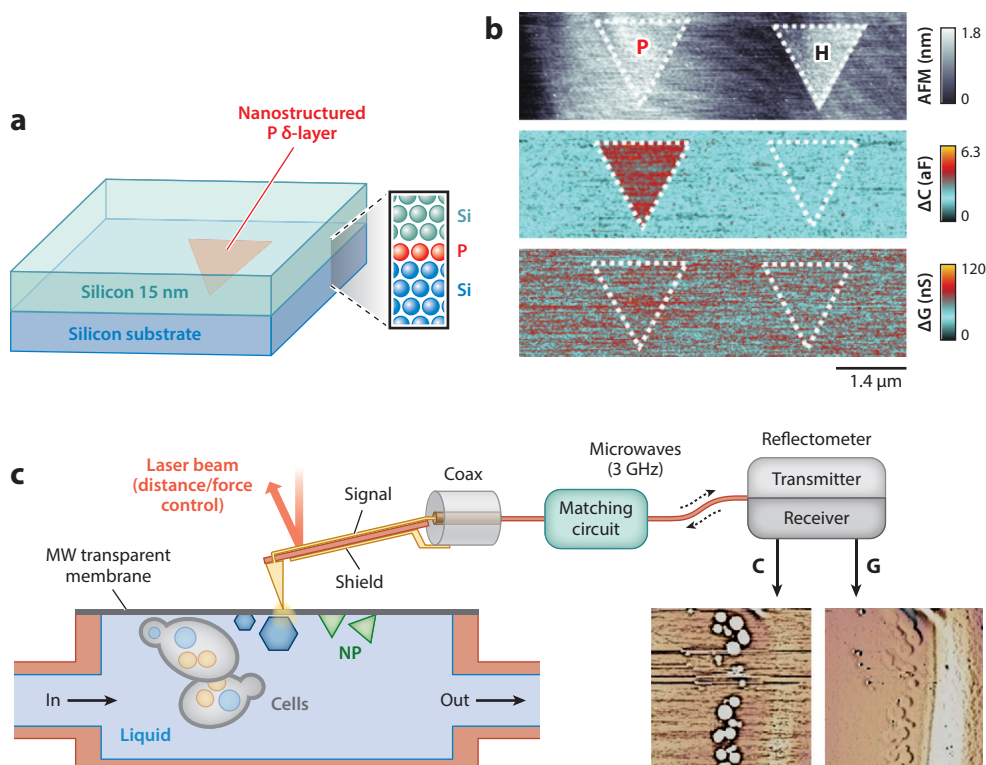




**Figure 3**

MIM imaging of a dielectric constant. (a) AFM (top), MIM-Im (bottom left), and MIM-Re (bottom right) images of few-layer  $\text{In}_2\text{Se}_3$  nanoflakes. The number of layers  $n$  is indicated in the AFM image. (b) Permittivity deduced from the MIM-Im data as a function of  $n$ . The blue circles and red squares represent two different batches of samples. (c) Schematic of microwave microscopy on a single bacterial cell. (d, top left) Capacitance image extracted from MIM-Im data. (Top right) 3D rendering of the topographic image. (Bottom) Comparison of numerically calculated intrinsic capacitance profiles for different dielectric constants (dashed lines) of the cell with the experimental intrinsic capacitance profile (solid line) after removing topographic cross-talk. The best agreement is found for  $\epsilon_r \sim 3$ –5. Abbreviations: AFM, atomic force microscopy; MIM, microwave impedance microscopy; MIM-Im, microwave impedance microscopy–imaginary part; MIM-Re, microwave impedance microscopy–real part. Panels a and b adapted with permission from Reference 38; copyright 2015 American Chemical Society (ACS). Panels c and d adapted with permission from Reference 42; copyright 2016 ACS.

**3.2.1. General-purpose studies.** With STM-assisted hydrogen resist lithography (45), it is now possible to pattern atomically thin P dopant nanostructures buried in Si wafers, as illustrated in **Figure 4a**. Characterization of these buried features is important for next-generation information processing, while at the same time presenting a major challenge to SPM tools that do not see through the surface. Using the MIM at  $f = 20$  GHz, Gramse et al. (46) were able to image a  $\delta$ -doped P layer encapsulated with 15-nm Si. In **Figure 4b**, the nanostructure incorporating P atoms is clearly seen in the capacitance image, whereas the control pattern with the hydrogen desorbed triangle on the right shows no measurable contrast. Note that the conductance channel displays little contrast for both the conductive P-region and the insulating H-region, consistent with the modeling in **Figure 2e**. The authors reported a lateral resolution of  $\sim 40$  nm and vertical



**Figure 4**

Examples of general-purpose conductivity imaging. (a) Schematic of the phosphorus  $\delta$ -doped layer buried in the Si substrate. (b, top to bottom) AFM image of a patterned phosphorus and de-passivated hydrogen layer and corresponding capacitance (from MIM-Im) and conductance (from MIM-Re) images. (c) Schematic layout of the experimental setup. The objects under study are enclosed in a capsule filled with a fluid (or gas) and separated from the probe by an ultrathin dielectric membrane transparent for MW radiation. Abbreviations: AFM, atomic force microscopy; MIM-Im, microwave impedance microscopy–imaginary part; MIM-Re, microwave impedance microscopy–real part; MW, microwave; NP, nanoparticle. Panels a and b adapted from Reference 46 (CC BY 4.0). Panels c and d adapted with permission from Reference 47; copyright 2016 American Chemical Society.

resolution of  $\sim 4$  nm, suggesting that the technique could aid the development of fabrication processes for surface-code quantum computers (46).

**Figure 4c** demonstrates the MIM imaging of objects in environmental cells (47). Here, the cells and metallic and insulating nanoparticles are enclosed in a capsule filled with a fluid and separated from the probe by an ultrathin dielectric membrane. The same configuration was also utilized to visualize the growth and stripping of Ag dendrites in liquid electrolytes, as well as live yeast cells immersed in glycerol beneath the  $\text{SiO}_2$  membrane (47). Since microwave imaging is free from radiolysis and radiation-induced damage, the approach can serve as an ideal framework for in situ studies of fragile organisms and research in reactive, toxic, or radioactive media. The technique can also be coupled with microfluidic platforms for electrophoretic and osmotic controls. The ease of operation and the compatibility with AFM platforms will continue to increase the popularity of MIM in various branches of materials research.



**3.2.2. Semiconductors.** Semiconductor devices, in which the electrical conductivity is controlled by doping, gating, temperature, or illumination, are the backbone of modern information technology. For the same reason, doped Si and GaAs devices have been the preferred samples for NSMM calibration (25, 28, 33, 48–50). Rather than dwelling on the traditional materials, however, we direct our attention to novel 2D and 1D semiconductors in this review.

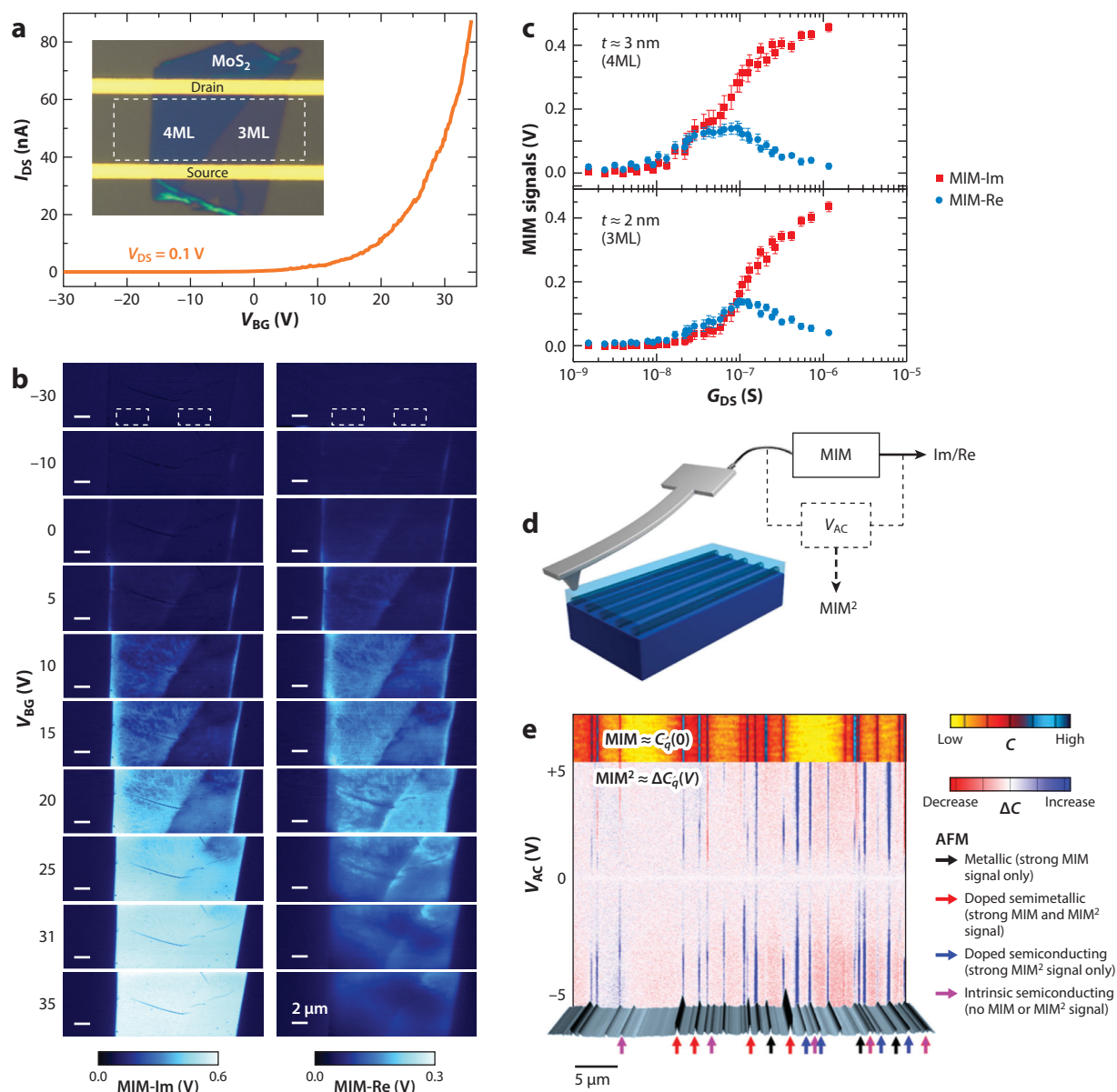
Semiconducting transition metal dichalcogenides, such as  $\text{MoS}_2$  and  $\text{WSe}_2$ , are in the lime-light of current materials research (51). These layered materials can be exfoliated into atomically thin 2D sheets with unique electrical and optical properties, which are attractive for nanoelectronics and optoelectronics. **Figure 5a** shows the transfer curve of a few-layer  $\text{MoS}_2$  flake in the field-effect transistor (FET) configuration (52). The device was covered by 15-nm-thick  $\text{Al}_2\text{O}_3$  to avoid direct contact between the metallic tip and the 2D sheet. **Figure 5b** displays selected MIM images within the channel region as a function of the back-gate voltage ( $V_{\text{BG}}$ ). As  $V_{\text{BG}}$  increased, the conductance signal emerged initially at the edges and then in the interior, with appreciable spatial nonuniformity (52). For quantitative analysis, average MIM-Im/Re signals on the four- and three-monolayer segments in **Figure 5b** are plotted in **Figure 5c**, which are consistent with the simulated response curves (**Figure 2e**). The results suggest that the contribution of defect-induced edge states to the total conductance is significant in the subthreshold regime but negligible once the bulk becomes conductive. The observation of conductance inhomogeneity also provides a guideline for future improvement of the device performance. Similar MIM works have been carried out on other 2D materials (53–62).

It is a common practice of MIM to modulate the local carrier concentration by applying a low- $f$  (kilohertz) voltage ( $V_{\text{AC}}$ ) on the tip (**Figure 5d**). The demodulated signal is recorded to form MIM-AC or microwave impedance modulation microscopy ( $\text{MIM}^2$ ) images during the scan (63). The configuration is similar to that in scanning capacitance microscopy, where the n- and p-doped regions exhibit opposite  $dC/dV$  signals (64). The combined MIM and  $\text{MIM}^2$  allows more detailed analysis of the sample. **Figure 5e** shows the MIM,  $\text{MIM}^2$ , and AFM data on an array of carbon nanotubes (CNTs) (63). For metallic CNTs, the large amount of carriers led to strong MIM and negligible  $dC/dV$  signals. At low  $V_{\text{AC}}$ , semiconducting CNTs displayed appreciable  $\text{MIM}^2$  signals, with p-doped tubes showing unipolar  $+\Delta C$  and intrinsic tubes showing bipolar  $\pm\Delta C$  on the two polarities of  $V_{\text{AC}}$ . Finally, for semimetallic CNTs, strong  $\text{MIM}^2$  signals appeared at high  $V_{\text{AC}}$ . The MIM tip modulation was also applied to other nanowire systems (65–67).

Using the FET configuration, either tip bias or sample bias, the Fermi level of the semiconducting materials can be locally modulated. The MIM/ $\text{MIM}^2$  is thus similar to the STM/scanning tunneling spectroscopy measurement, with lower resolution but much relaxed requirements for sample preparation and environment control. MIM also would be useful for novel semiconductor devices, such as ferroelectric FETs (68) and electric double-layer transistors (69), which was demonstrated recently (70, 71).

**3.2.3. Photoconductors.** Photoconductivity, the drastically altered electrical conductivity in a material under optical illumination, is widely exploited in optoelectronics. With some exceptions for semimetals and correlated materials, most photoconductors are also semiconductors. Considering the addition of a light source and optical components in the setup, however, light-illuminated microwave impedance microscopy (iMIM) deserves a separate section here. Note that iMIM, which measures the intrinsic photoconductivity without the need of electrodes, is fundamentally different from scanning photocurrent microscopy, which maps out the photo-generated current across the source and drain electrodes when a focused laser beam scans over the device (72).

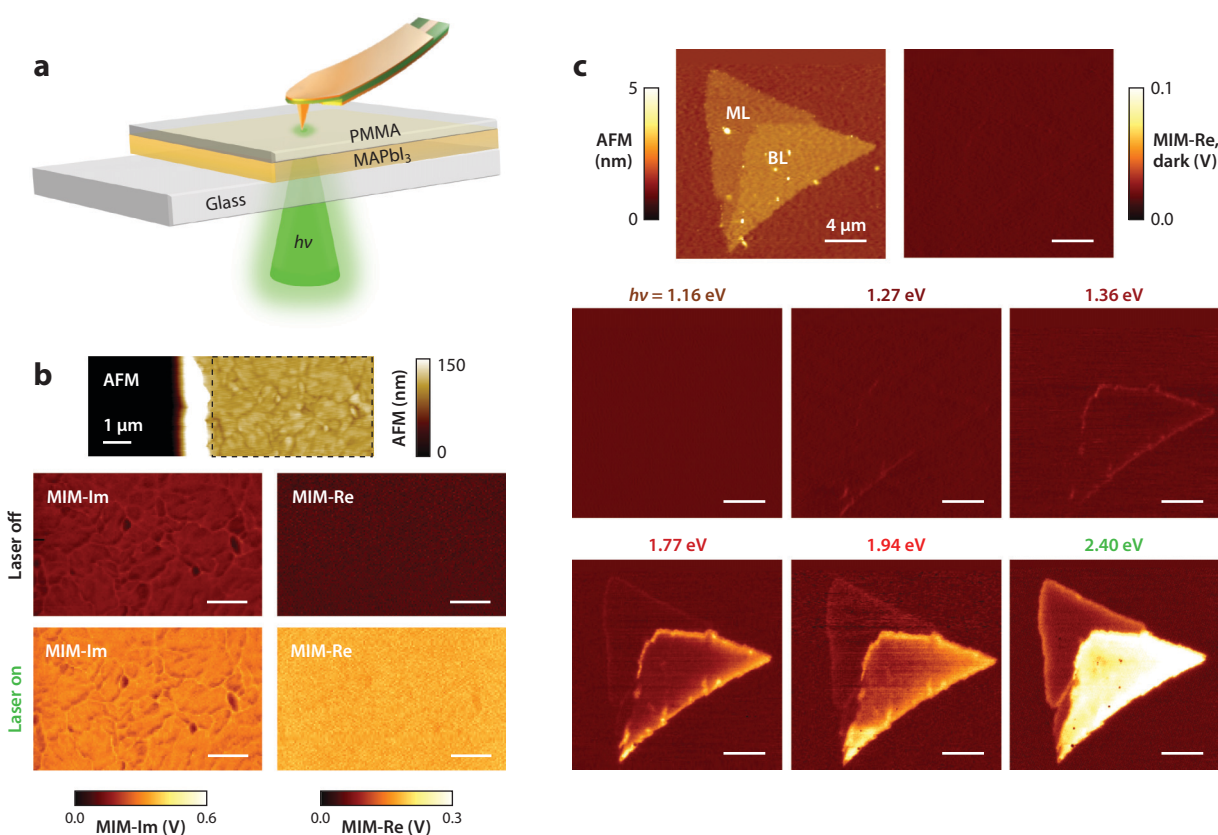
The worldwide surge of research interest in organic-inorganic trihalide perovskites, e.g.,  $\text{CH}_3\text{NH}_3\text{PbI}_3$  or  $\text{MAPbI}_3$ , has led to a phenomenal increase in the power conversion efficiency from 3.8 to 24.2% in the past few years (73). **Figure 6a** shows a schematic of a recent iMIM study, in which the tip was aligned with a focused laser beam illuminated through a transparent glass substrate (74). When the perovskite solar cell film was in the dark, the MIM signals indicated the absence of mobile carriers. When illuminated by an above-gap laser with  $\lambda = 532$  nm, however, clear photoconductivity was seen in the MIM images (**Figure 6b**). Interestingly, while the MIM-Im image shows appreciable topographic cross-talk, the MIM-Re image is rather uniform across



(Caption appears on following page)

**Figure 5** (Figure appears on preceding page)

MIM imaging on semiconductors. (a) Transfer characteristics of the back-gated MoS<sub>2</sub> transistor. The inset shows a picture of the device. (b) Selected MIM images of the sample in the dashed box in panel a. (c) Averaged MIM signals as a function of the source-drain conductance for the 4ML and 3ML regions in the dashed boxes in panel b. (d) Schematic of MIM and MIM<sup>2</sup> experiments on encapsulated carbon nanotubes. (e, top to bottom) MIM, MIM<sup>2</sup>, and AFM topography scans of the same nanotube array, illustrating the complementary nanoscale data obtainable with each technique. The signatures have been categorized for the four electronic nanotube types indicated by the colored arrows: strong MIM signal only (black, metallic), strong MIM<sup>2</sup> signal only (blue, lightly doped semiconducting), both strong MIM and MIM<sup>2</sup> signals (red, semimetallic), and no signal with either technique (purple, intrinsic semiconducting). Abbreviations: 3ML, three monolayer; 4ML, four monolayer; AFM, atomic force microscopy; MIM, microwave impedance microscopy; MIM<sup>2</sup>, microwave impedance modulation microscopy; MIM-Im, imaginary microwave impedance microscopy; MIM-Re, real microwave impedance microscopy. Panels a–c adapted with permission from Reference 52; copyright 2016 Proceedings of the National Academy of Sciences. Panels d and e adapted with permission from Reference 63; copyright 2016 American Chemical Society.

**Figure 6**

Photoconductivity imaging by iMIM. (a) Schematic of the hybrid perovskite sample and the iMIM setup with bottom illumination through the transparent glass substrate. Scanning is accomplished by moving the sample stage while fixing the laser beam and the probe tip, which are aligned before the experiment. (b) AFM (top) and MIM-Im/Re images (inside the black dashed rectangles in the AFM data) when the 532-nm laser is turned off (middle) and on (bottom). (c) AFM and MIM-Re images of an ML-BL WSe<sub>2</sub> flake when illuminated by diode lasers with different energies, showing the sequential emergence of photoconductivity in different regions of the sample. Abbreviations: AFM, atomic force microscopy; BL, bilayer; iMIM, light-illuminated microwave impedance microscopy; MIM-Im, microwave impedance microscopy–imaginary part; MIM-Re, microwave impedance microscopy–real part; ML, monolayer; PMMA, polymethyl methacrylate. Panels a and b adapted from Reference 74 (CC BY 4.0). Panel c adapted with permission from Reference 77; copyright 2018 American Chemical Society.

grains and grain boundaries, suggesting that microstructures do not lead to strong spatial variations of the intrinsic photo-response. The same work also showed that the degradation process began with the disintegration of grains rather than nucleation and propagation from visible grain boundaries (74). Such understanding of the defect structures, grain boundaries, and air stability is important to improve the device performance towards commercialization.

Vertical and lateral heterostructures of 2D materials provide tremendous flexibility to engineer local band structures (75, 76). **Figure 6c** shows the MIM images of photoconductivity in a monolayer-bilayer  $\text{WSe}_2$  lateral heterostructure under multiple excitation lasers (77). As the photon energy increased from 1.16 to 2.40 eV, the light-induced conductivity first appeared along the hetero-interface and bilayer edge, then along the monolayer edge, inside the bilayer area, and finally in the interior of the monolayer region. The sequential emergence of mobile carriers in different sections of the sample can be used to construct a qualitative configuration of local energy gaps, which is consistent with the density functional theory calculations (77). The results suggest the exciting possibility of band-gap engineering on the length scale of nanometers.

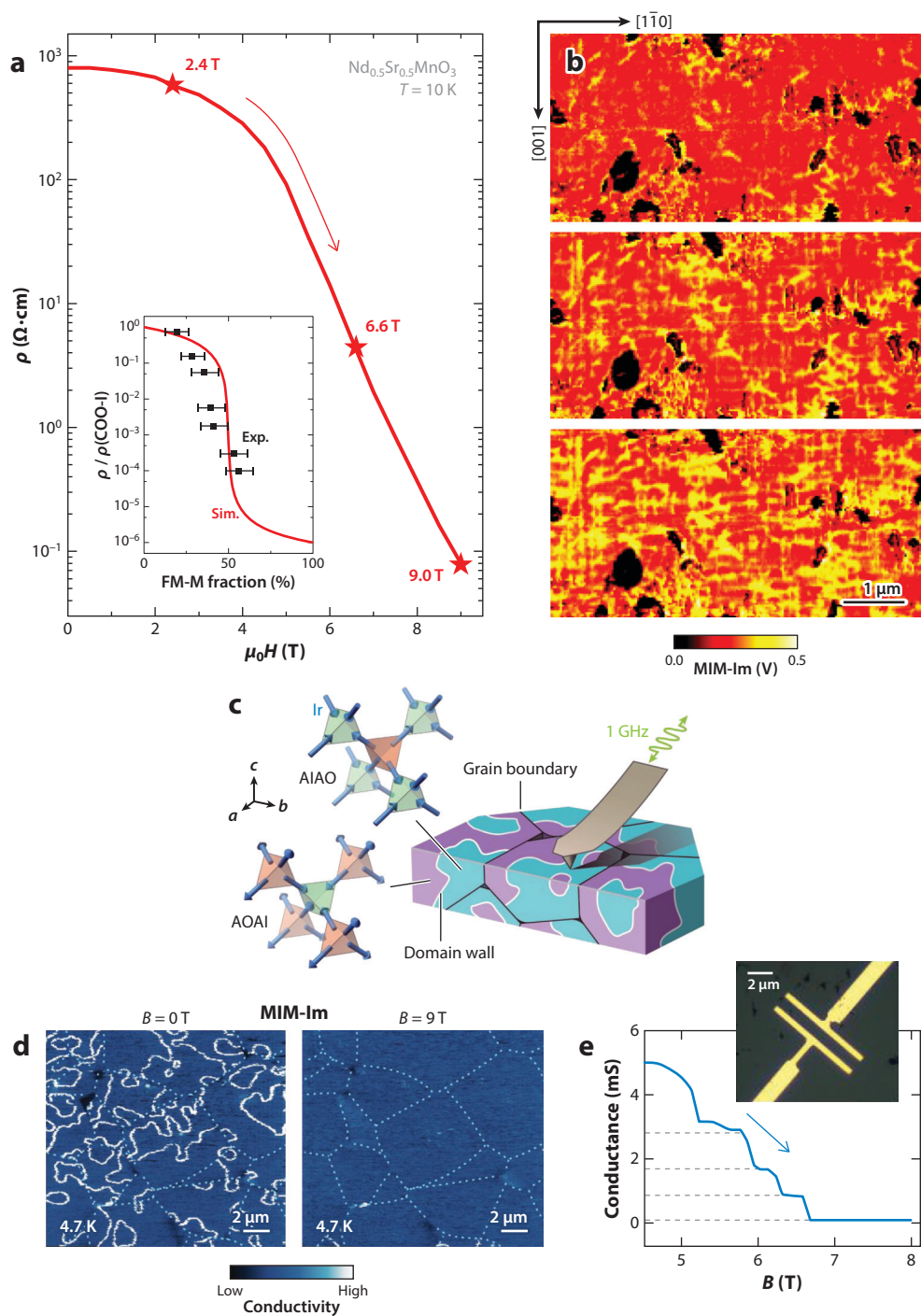
Photoconductivity imaging is still in its infancy (74, 77, 78–80) and will likely see rapid growth in the next few years. First, it is straightforward to use a broadband light source to determine the onset of free-carrier generation. The electronic gap measured by iMIM is different from the optical gap determined from absorption experiments, as the latter is reduced by the exciton binding energy (51). Second, by moving the tip with respect to the laser spot, a diffusion map of the photo-generated carriers can be obtained. Third, by using a pulsed laser and fast electronics, the microwave response can be temporally resolved down to 1–10 ns, which will be a powerful method to study the dynamics of photo-excited electrons. Finally, the measurements can be incorporated in a cryogenic environment to suppress the thermal excitation.

**3.2.4. Strongly correlated systems.** Strongly correlated materials are a wide class of compounds whose behaviors cannot be described by noninteracting or weakly interacting physics (81). A defining feature of these systems is the emergence of unusual electronic and magnetic properties (82). In particular, the electron correlation may lead to the coexistence of competing states with drastically different electronic structures, whose spatial configurations closely reflect the underlying physics. MIM would be an ideal tool to probe such coexisting domains or domain boundaries.

**Figure 7a** shows the low- $T$  magneto-transport data of the colossal magnetoresistive material  $\text{Nd}_{0.5}\text{Sr}_{0.5}\text{MnO}_3$  with coexisting charge/orbital-ordered insulating phase and ferromagnetic metallic (FM-M) phase (83). As the magnetic field increased from 2.4 to 9.0 T, the macroscopic transport resistivity dropped by four orders of magnitude, accompanied by substantial expansion of the FM-M phase in the MIM-Im image (**Figure 7b**). The FM-M phase at high field formed an interconnected network, with a volume fraction consistent with the 2D percolation transition (inset of **Figure 7a**). Strikingly, the prominent FM-M filaments aligned preferentially along the [001] and  $[1\bar{1}0]$  directions of the substrate, suggesting the anisotropic elastic strain as the key interaction here (83). Metal-insulator transitions in complex oxides such as  $\text{VO}_2$  (84, 85) and  $\text{Pr}_{1-x}(\text{Ca,Sr})_x\text{MnO}_3$  (86) were also imaged by MIM.

In a magnetic insulator, whether metallic behavior occurs at the DWs is a long-standing question that remains elusive experimentally (87). **Figure 7c** shows a schematic of the low- $T$  insulator  $\text{Nd}_2\text{Ir}_2\text{O}_7$  with unusual all-in-all-out (AIAO) magnetic orders (88). The zero-field MIM-Im image of a polished polycrystalline sample is displayed in **Figure 7d**, showing highly conductive curvilinear features with respect to the insulating bulk (89). In contrast, most conductive curves disappeared under 9 T, which aligned the magnetic domains and eliminated the AIAO DWs. As supporting evidence, the transport conductance across two electrodes fabricated on the sample





(Caption appears on following page)

**Figure 7** (Figure appears on preceding page)

MIM application on strongly correlated materials. (a) Resistivity of an  $\text{Nd}_{0.5}\text{Sr}_{0.5}\text{MnO}_3$  thin film during the field sweep at 10 K. The three fields at which MIM images were taken are marked by red stars. The inset shows a comparison between the 2D square-lattice simulation (red curve) and the experimental data (black squares). (b) MIM-Im images taken at 2.4, 6.6, and 9.0 T. The ferromagnetic metallic phase (yellow) exhibits higher signals than the charge/orbital-ordered insulating phase (red). The black particles are  $\text{MnO}_x$  defects. (c) Illustration of the MIM scanning setup on polished polycrystal  $\text{Nd}_2\text{Ir}_2\text{O}_7$ , showing the spin configuration of the AIAO order and its variations. Domain walls can exist between the two variations. (d, left) MIM-Im image taken after zero-field cooling from 40 to 4.7 K. The dotted lines are grain boundaries, and the dark spots are voids between grains. Curvilinear features much more conductive than the bulk are AIAO magnetic domain walls. (Right) MIM-Im image in the same region after a thermal cycle to 40 K and cooling back to 4.7 K in 9 T. Most grains become single domain. (e) Typical conductance versus magnetic field of a microelectrode device (inset) at 2 K, showing staircase-like drops. Abbreviations: AIAO, all-in-all-out; AOAI, all-out-all-in; exp., experiment; FM-M, ferromagnetic metallic; MIM, microwave impedance microscopy; MIM-Im, microwave impedance microscopy–imaginary part; sim. simulation. Panels a and b adapted with permission from Reference 83; copyright 2010 American Association for the Advancement of Science (AAAS). Panels c–e adapted with permission from Reference 89; copyright 2015 AAAS.

showed abrupt drops as the field ramped up (**Figure 7e**), consistent with the sequential removal of DWs between the electrodes. The results unambiguously establish the metallic nature of DWs in this AIAO magnetic insulator. Enhanced conductivity was also observed in charge-ordered DWs in the layered manganite  $\text{Pr}(\text{Sr}_{0.1}\text{Ca}_{0.9})_2\text{Mn}_2\text{O}_7$  (90).

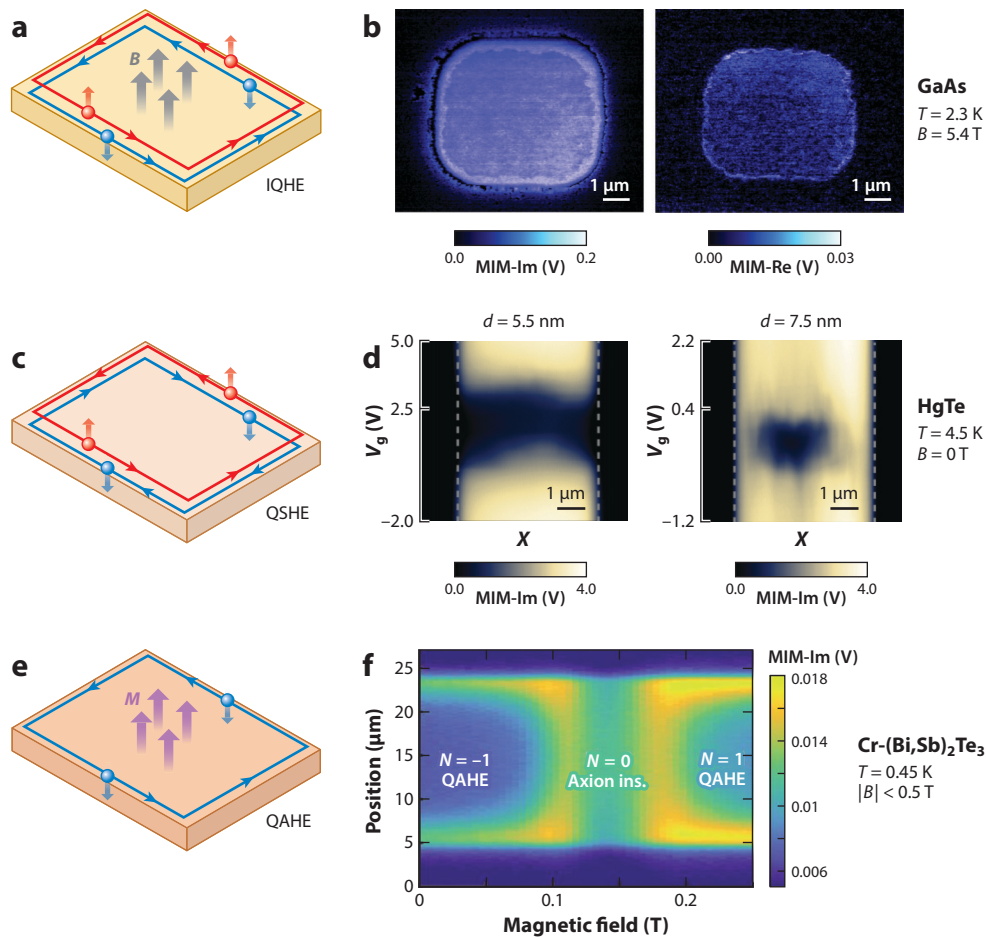
Local imaging on strongly correlated materials has been very fruitful to date, as exemplified by the study of nanoscale inhomogeneity in unconventional superconductors by STM (91). The highly resistive phases in many complex oxides, however, are difficult to study by a tunneling experiment. Similarly, while NSOM has become increasingly important in the imaging of correlated systems (92), it is mostly sensitive to the optical contrast between metals and insulators. As already demonstrated above, MIM is in a unique position to continue the investigation on these intriguing quantum materials.

**3.2.5. Topological quantum systems.** The research on the topologically ordered phase of matter (93, 94) has been in the central stage of quantum science and engineering. In particular, the dissipationless edge or surface states may enable novel memory and logic technologies with minimal energy consumption. While these conductive channels have been demonstrated by macroscopic measurements, much remains to be explored on their microscopic details.

The integer quantum Hall effect (IQHE) is characterized by the exact quantization of the Hall resistance and vanishing longitudinal resistance (95). As illustrated in **Figure 8a**, the topological boundary states are formed by the intersection between Landau levels (LLs) and the Fermi level ( $E_F$ ) of a high-mobility 2D electron system (2DES) under high magnetic ( $B$ ) fields. **Figure 8b** shows the MIM images at  $T = 2.3$  K and  $B = 5.4$  T of a GaAs/AlGaAs 2DES etched into isolated disks (96). Immediately inside the physical boundary, the strong MIM-Im and zero MIM-Re signals are indicative of a metallic strip around the perimeter. The MIM imaging also enabled quantitative measurement of the width of edge states, and the result is in good agreement with the electrostatic model (97). A follow-up experiment was performed in a graphene flake, and the IQHE edges were again imaged (98).

Different from the IQHE at high  $B$ -fields, the quantum spin Hall effect (QSHE) shows a quantized spin Hall conductance and a vanishing charge Hall conductance at  $B = 0$  (99, 100). As illustrated in **Figure 8c**, the QSHE features a bulk energy gap and a pair of gapless helical edge states carrying opposite spins. **Figure 8d** shows MIM line scans across two  $\text{HgTe}/(\text{Hg}_{0.3}\text{Cd}_{0.7})\text{Te}$  quantum well (QW) devices as  $E_F$  was tuned by the back gate (101). For the topologically trivial





**Figure 8**

Microwave imaging of topological edge states. (a) Schematic of the IQHE under high magnetic fields. (b) MIM images at  $B = 5.4$  T and  $T = 2.3$  K of a high-mobility GaAs 2D electron system. Conductive edge channels are clearly seen around the physical boundary of the etched disk. (c) Schematic of the QSHE under zero magnetic field. (d) Zero-field MIM single-line scan versus gate voltage across an HgTe/(Hg<sub>0.3</sub>Cd<sub>0.7</sub>)Te device with a quantum-well thickness of 5.5 nm (left) and 7.5 nm (right). Within the bulk gap region, conductive edges are absent in the topologically trivial sample on the left and present in the QSHE sample on the right. (e) Schematic of the QAHE with an out-of-plane magnetization. (f) MIM-Im line scan across a Cr-doped (Bi,Sb)<sub>2</sub>Te<sub>3</sub> sample versus the magnetic field, showing the evolution of edge states across the two QAHE states and the axion insulator state. Abbreviations: axion ins., axion insulator; IQHE, integer quantum Hall effect; MIM, microwave impedance microscopy; MIM-Im, microwave impedance microscopy–imaginary part; MIM-Re, microwave impedance microscopy–real part; QAHE, quantum anomalous Hall effect; QSHE, quantum spin Hall effect. Panel *b* adapted with permission from Reference 96; copyright 2011 American Physical Society. Panel *d* adapted from Reference 101 (CC BY 4.0). Panel *f* adapted with permission from Reference 107; copyright 2019 Proceedings of the National Academy of Sciences.

sample with a QW thickness  $d$  of 5.5 nm (below the critical value of 6.5 nm), the entire sample became insulating inside the bulk gap. For the nontrivial sample with  $d = 7.5$  nm, however, the bulk turned insulating but the two edges remained conductive as  $E_F$  passed through the bulk gap, consistent with the QSHE physics in HgTe (102, 103). The QSHE edge states in monolayer WTe<sub>2</sub> flakes were also visualized by MIM in a recent work (104).

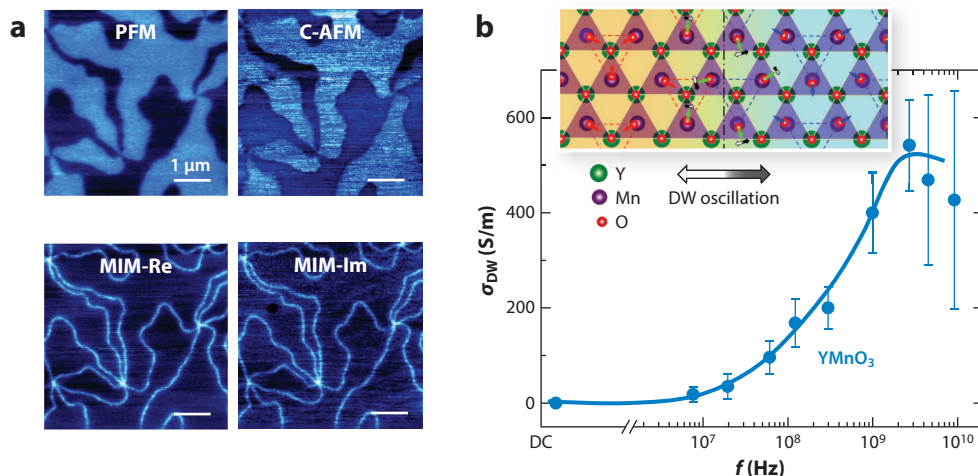
In analogy to the  $N = 1$  ( $N$  being the LL index or the first Chern number) IQHE state, the quantum anomalous Hall effect (QAHE) is characterized by a chiral 1D edge current that encloses an insulating bulk (105, 106). The time-reversal symmetry, however, is broken by a spontaneous out-of-plane magnetization (**Figure 8e**) rather than an external  $B$ -field. **Figure 8f** shows the MIM line scans across a Cr-(Bi,Sb)<sub>2</sub>Te<sub>3</sub> sample with two magnetically doped layers (107), whose magnetization was controlled by a small  $B$ -field ( $< 0.5$  T). For the  $N = \pm 1$  states, the hallmark of QAHE, i.e., dissipationless edges and insulating bulk, was vividly imaged by the MIM. In between the two QAHE states, however, the system entered a topologically distinct axion insulator ( $N = 0$ ) state (108–110) with spatially uniform conductivity, where the magnetization direction was opposite in the two magnetic layers. The visualization of both QAHE and axion insulator states further testifies to the ability of MIM to probe exotic quantum states.

In the past decade, many topologically nontrivial states have been theoretically proposed and experimentally realized. Given its nanoscale resolution, strong electrical contrast, noninvasive nature, and compatibility with a high- $B$ /low- $T$  environment, MIM may become indispensable for probing topological materials. In the meantime, the interpretation of MIM data in the quantum regime needs to be improved to include ballistic transport (rather than diffusive transport in **Figure 1d**) for high-mobility samples and topological channels, as well as the full resistivity tensor with a nontrivial Hall contribution. It is likely that what we have seen here is merely the tip of the iceberg, leaving open space for future explorations.

### 3.3. Imaging Dielectric Loss

The dielectric loss in a material quantifies the dissipation of EM energy due to dipolar relaxation or the excitation of collective modes (111). At a fixed frequency, the dipolar loss  $\epsilon''\omega$  is indistinguishable from the electronic conduction  $\sigma$  at the circuit level. To separate the contributions between mobile carriers and bound charges, therefore, one must perform simultaneously DC (usually by C-AFM) and AC (usually by  $f$ -dependent MIM) microscopy for the mapping of local dielectric loss in complex systems.

When a multidomain ferroelectric is placed in microwave fields, the coupling between the spontaneous polarization and the alternating  $E$ -field may lead to periodic oscillation of the DWs and, thus, dielectric loss (112–114). On the other hand, it is now widely accepted that many ferroelectric DWs can host electrical conduction due to the presence of free carriers (115, 116). For weakly charged walls in KNbO<sub>3</sub> crystals (117) and transiently formed charged walls in lead zirconate titanate thin films (118), the gigahertz DW conductivity measured by MIM is indeed dominated by charge carriers. The situation, however, is different for the neutral walls on hexagonal manganites (h-RMnO<sub>3</sub>). **Figure 9a** shows piezo-force microscopy (PFM), C-AFM, and MIM images on the [001] surface of YMnO<sub>3</sub> (119). The C-AFM data showed that the DWs are more resistive than the adjacent domains (120) with a DC conductivity  $\sim 10^{-3}$  S/m. On the other hand, the MIM images in the same area indicate very strong gigahertz response at the DWs. For comparison, the DW signals were quantified by an effective gigahertz conductivity, which is as high as  $\sim 400$  S/m from the FEA. The  $f$ -dependent DW conductivity (**Figure 9b**) was measured by a series of MIM electronics, showing a peak-like feature around 3 GHz. The result established that the DW vibration (inset of **Figure 9b**), rather than electronic conduction, is responsible for the



**Figure 9**

Imaging of dielectric loss by MIM. (a) PFM, C-AFM, and MIM images acquired on the [001] surface of YMnO<sub>3</sub>. The cloverleaf-like domain patterns are seen in the PFM and C-AFM images, whereas the DWs are highlighted in the MIM data. (b) Frequency-dependent effective DW conductivity. The blue line is a visual guide. The inset shows an atomistic view of YMnO<sub>3</sub> in the [001] plane across the interlocked antiphase boundary and ferroelectric DW. The black and white double-headed arrow illustrates the amplitudes and directions of the periodic DW sliding. Abbreviations: C-AFM, conductive atomic force microscopy; DW, domain wall; MIM, microwave impedance microscopy; MIM-Im, microwave impedance microscopy-imaginary part; MIM-Re, microwave impedance microscopy-real part; PFM, piezo-force microscopy. Figure adapted from Reference 119 (CC BY 4.0).

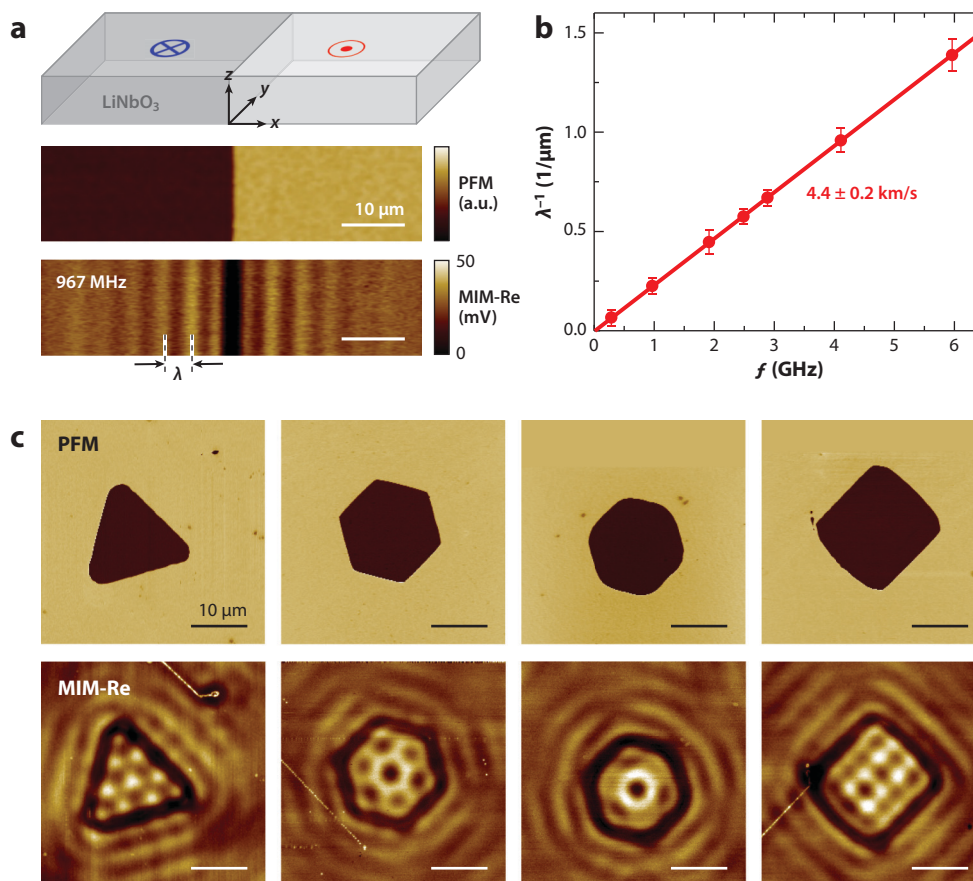
pronounced dielectric loss (119). Similar behavior was also seen in hexagonal ferrite (h-RFeO<sub>3</sub>) (121). It should be emphasized that the DW physics in ferroelectric YMnO<sub>3</sub> is different from that in the magnetic insulator Nd<sub>2</sub>Ir<sub>2</sub>O<sub>7</sub> (Figure 7d) (89), where the walls are equally conductive at DC and microwave frequencies.

The measurement of an  $f$ -dependent dielectric response will undoubtedly lead to follow-up investigations on other ferroelectric DWs. More importantly, broadband MIM will open up enormous opportunities to locally probe materials with gigahertz collective modes, such as charge density waves (122) and pinned Wigner crystals in 2DES (123). Since the dielectric loss in these samples is specific to the microwave frequency, MIM would be the only technique to map out their gigahertz dynamics in the nanoscale.

### 3.4. Imaging Electroacoustic Transduction

In a piezoelectric material, the mechanical stress is coupled to the electric potential by the linear electromechanical interaction. An oscillating gigahertz  $E$ -field, therefore, will launch acoustic waves in the sample. Such electroacoustic transduction has found many applications in electronic components such as surface acoustic wave (SAW) devices (124). For MIM on piezoelectrics, this new channel of energy dissipation, in addition to the usual dielectric response, should be taken into account for image analysis (125, 126).

Figure 10a shows the results on a  $z$ -cut LiNbO<sub>3</sub> sample with one long DW (126). The two oppositely polarized domains are seen in the PFM image. The MIM-Re image, on the other hand, exhibits clear interference fringes around the wall. As the electrical conductance of LiNbO<sub>3</sub>



**Figure 10**

MIM imaging on piezoelectric power transduction. (a, top to bottom) Schematic, PFM, and MIM-Re images of the *z*-cut LiNbO<sub>3</sub> crystal with a single domain wall in the middle. The MIM image exhibits clear interference fringes, with a period of  $\lambda$  labeled in the plot, around the DW. (b) Linear relation between  $1/\lambda$  and the MIM frequency. The slope corresponds to a wave velocity of  $4.4 \pm 0.2$  km/s. (c) PFM and MIM-Re images of four LiNbO<sub>3</sub> coral domains. Clear interference patterns due to the superposition of ripples around each DW are seen in the MIM-Re data. Abbreviations: DW, domain wall; MIM, microwave impedance microscopy; MIM-Re, microwave impedance microscopy–real part; PFM, piezo-force microscopy. Figure adapted with permission from Reference 126; copyright 2018 Proceedings of the National Academy of Sciences.

is negligible, the MIM-Re contrast indicates that the microwave energy is dissipated through piezoelectric transduction rather than Ohmic loss. In **Figure 10b**, the measured  $1/\lambda$  ( $\lambda$  being the period of the MIM-Re fringes) is plotted as a function of the MIM frequency. The slope of  $4.4 \pm 0.2$  km/s closely matches that of the *x*-propagating pseudo-SAW on *z*-cut LiNbO<sub>3</sub> (127). Using electrometrical FEA, one can simulate the one- $\lambda$  oscillation in energy loss as the tip moves away from the DW, which stems from the nonlocal interaction between electric fields and elastic waves (126). The interference of piezoelectric transduction can be further appreciated in LiNbO<sub>3</sub> coral domains (126), as seen from **Figure 10c**.

MIM of ferroelectric domains reveals certain internal degrees of freedom in piezoelectric and elastic tensors, which are not accessible by measurements of the acoustic displacement

fields. Continued work along this line may open a new research frontier to explore nanoscale electroacoustic phenomena in complex structures.

### 3.5. Beyond Electrical Impedance Imaging

While the majority of NSMM applications to date are the electrical impedance type, exceptions exist for imaging other material properties, which are briefly discussed here.

**3.5.1. Imaging magnetic impedance.** To probe the microwave magnetic response, one needs to modify the lumped-element model in **Figure 1a** to include the tip-sample mutual inductance. The coupling to the electrically open tip described above is relatively weak (128). On the other hand, an electrically short or loop tip that maximizes microwave magnetic fields around the probe head would enable study of the local permeability and magnetic resonance (6). Given the difficulty of forming a nanoscale loop probe and the competing technology in magnetic imaging, the success of this research area is still pending future developments.

**3.5.2. Imaging gigahertz nonlinearity.** The local higher-order dielectric or magnetic properties can be measured by probing the nanoscale nonlinear response. In practice, one can either apply kilohertz modulation to the gigahertz excitation and demodulate the higher-order signals or directly detect the higher harmonics of the gigahertz signals. By using the former scheme, scanning nonlinear dielectric microscopy has achieved appreciable success in ferroelectrics (129) and semiconductors (130). Similarly, by using a loop probe and the latter scheme, investigators have also detected the nonlinear Meissner response at edges and defects in superconducting samples (131).

**3.5.3. Imaging microwave fields.** Near-field microwave probes have been used to map out the gigahertz fields in microwave resonators (132) and metamaterials (133), where the characteristic length scale is determined by the EM wavelength and mesoscopic resolution is not necessary. The spatial mapping of a time-varying  $E$ -field that accompanies SAWs in piezoelectrics, however, does require a nanoscale probe due to the micrometer-sized gigahertz acoustic wavelength. The configuration is different from the imaging of transduction loss (Section 3.4) in that the tip is acting as a receiver to detect the microwave fields launched from a nearby transducer. Such acoustic field imaging has been recently demonstrated on SAW components (134) and resonators (135), which may find applications in electroacoustic devices, quantum acoustics, and phononic crystals.

## 4. SUMMARY AND OUTLOOK

In summary, the past decade has witnessed the upturn of near-field microwave microscopy, especially in the form of MIM, in various branches of materials research. By quantitatively imaging on the nanoscale, the dielectric constant and conductivity can be obtained on novel semiconductors, photoconductors, and biological materials. Implementation of the technique in cryogenic and magnetic-field environments has led to important discoveries in materials with strong correlation and topological order. The understanding of MIM contrast in ferroelectric DWs and piezoelectrics also expands its territory into dipolar materials and electromechanical systems.

Having surveyed the major progress of MIM in recent years, we conclude that the technique has arrived at a critical juncture for future prosperity. The success of a particular SPM mode depends on multiple conditions: (a) commercial companies supplying the specialized probes, electronics, and platforms compatible with the module; (b) a broad user community, ideally including both



industrial teams and academic groups; and (c) widespread applications, ideally including general-purpose imaging and killer applications in frontier research. As detailed in this review, the rapid progress of MIM in the past decade has addressed these key points with positive answers, which certainly bodes well for the future.

It is also interesting to ponder the ultimate limits of microwave microscopy in the next few decades. A number of desired functionalities and applications are discussed below.

#### 4.1. Spatial Resolution

Except for the configurations based on tunneling (136) and nonlinearity (137), the spatial resolution of NSMM with linear dielectric contrast has been limited to  $\sim 50$  nm for most reports to date. Further improvement would require a concerted effort to construct durable and sharp tips, sensitive electronics, and reliable feedback control from the scanner. We anticipate that sub-10-nm resolution can be achieved within the next decade.

#### 4.2. Operation Frequency

Beyond the current MIM frequency band (from low MHz to  $\sim 20$  GHz), the probe and electronics become progressively harder to construct. On the other hand, material properties in the millimeter-wave (30–300 GHz) regime are of great interest for civil and military applications. Microscopy around  $10^{11}$  Hz will also bridge the gap between the current frequencies of NSMM (up to  $\sim 10^{10}$  Hz) and NSOM (down to  $\sim 10^{12}$  Hz). With efforts along this direction (138–140), near-field imaging throughout the EM spectrum may happen in one or two decades.

#### 4.3. Industrial Applications

As discussed above, microwave microscopy could be appealing for multi-billion-dollar industries such as semiconductor technology, photovoltaics, and SAW devices. Industrial applications usually require high-throughput, large-scale (e.g., wafer-scale) inspection and fast (e.g., video-speed) operation, which necessitate building suitable probes, electronics, and scanning platforms. The ability to zoom out for coarse imaging and zoom in for fine scanning would be desirable as well. It is hopeful that MIM will become the standard tool for multiple industries in the future.

#### 4.4. Academic Applications

The broad range of MIM applications reviewed here is a good start for cutting-edge scientific studies. For complex systems with various degrees of freedom, it is usually advantageous to perform multimodality microscopy. One may further envision the integration of MIM with NSOM, electrostatic force microscopy, Kelvin probe force microscopy, and magnetic force microscopy, among others, in one experiment. In addition, the research of quantum materials may require extreme conditions, such as milli-Kelvin temperature in a dilution refrigerator (141) or ultrahigh vacuum in a molecular-beam epitaxy chamber. Efforts are underway to incorporate MIM into these instruments, which will continue to bring us exciting discoveries in materials research.

### DISCLOSURE STATEMENT

K.L. holds a patent on MIM technology, which is licensed to PrimeNano Inc. for commercial instruments. The terms of this arrangement have been reviewed and approved by the University of



Texas at Austin in accordance with its policy on objectivity in research. The remaining authors are not aware of any affiliations, memberships, funding, or financial holdings that might be perceived as affecting the objectivity of this review.

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