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Scanning nanoscale multiprobes for conductivity measurements

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We report fabrication and measurements with two- and four-point probes with nanoscale dimensions, for high spatial resolution conductivity measurements on surfaces and thin films. By combination of conventional microfabrication and additive three-dimensional nanolithography, we have obtained electrode spacings down to 200 nm. At the tips of four silicon oxide microcantilevers, narrow carbon tips are grown in converging directions and subsequently coated with a conducting layer. The probe is placed in contact with a conducting surface, whereby the electrode resistance can be determined. The nanoelectrodes withstand considerable contact force before breaking. The probe offers a unique possibility to position the voltage sensors, as well as the source and drain electrodes in areas of nanoscale dimensions. © 2000 American Institute of Physics.

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I. INTRODUCTION

Four-point measurements have played an important role in understanding the electrical properties of solid state bulk materials and films for many decades.¹ A typical four-point configuration is a linear array of four equidistant electrodes, with two outer electrodes performing as source and drain, while the voltage difference is measured across the inner electrodes. The conductivity can be derived from the voltage-to-current ratio, while compensating for geometrical effects related to the thickness, size, and shape of the sample. Due to the electrical separation of the source-drain electrodes and the voltage electrodes, the conductivity is measured essentially without contact resistance. Common types of four-point probes are macroscopic spring-loaded electrodes, that can be positioned freely on a surface,² and smaller, lithographically fabricated in-plane electrodes fixed on the surface.³ The macroprobes are in many cases inconvenient because of their size and the large contact forces exerted on the samples, while the in-plane electrodes cannot be repositioned. The four-point technique is suited where the conductivity varies slowly on the scale of the electrode spacing.⁴ In order to measure systems that exhibit conductivity variations on a small scale, it is preferable to reduce the electrode spacing below the relevant scale, yet maintaining the ability to reposition them. Later we describe a solution to this challenge.

II. PROBE FABRICATION

Using conventional photolithographic microprocessing techniques, we have fabricated four-point probes consisting of four soft and flexible metallized SiO₂ microcantilevers,⁴ with electrode spacings down to 1.5 μm, as shown in Fig. 1(b). To reduce the gap even further, we take advantage of

electron-beam induced deposition, which is a constructive three-dimensional nanolithography technique.^{5–10} By focusing the electron beam of a scanning electron microscope on the ends of the microcantilevers, hydrocarbon molecules present in small concentrations near the beam spot are cracked. This leads to the formation of a narrow rod of carbon residues, growing in the direction of the beam. Lengths of several microns are easily obtained by continuation of the process, and such tips have successfully been used for high aspect ratio atomic force microscopy tips,⁵ nanoparticle manipulation,⁶ and nanolithography.^{7,8}

To avoid charging of the oxide cantilevers by the electron beam, and thus a poor growth rate and tip quality,⁷ the microcantilevers are coated with a thin metallic layer (100 Å Ti/800 Å Au) prior to tip deposition. By keeping the electrodes at the ground potential of the scanning electron microscope (SEM), a drain for the electron beam is provided. In order to make the nanotips conducting the nanotips are metallized a second time (100 Å Ti/600 Å Au). An alternative approach could be to grow the tips in the presence of controlled amounts of metallo-organic compounds inside the vacuum chamber.^{9,10}

We used a JEOL 6340F field emission microscope operating at a base pressure of 10⁻⁸ Torr. With an acceleration voltage of 10 kV and beam currents in the range 3–6 pA we obtained a growth rate of 250 nm/min. The beam current is comparable to the 1.9 pA found to be optimal by Wendel *et al.*⁷ in terms of growth rate and resulting sharpness of the tips. Similarly to these findings,⁷ we observe a decay of the growth rate as the tip grows longer.

The carbon tips turn out to be mechanically strong and durable, as demonstrated in earlier works.^{6,7} It has been suggested that the tip material is diamond-like, without being brittle, which would explain the exceptional durability.⁷

By tilting the microprobe with respect to the beam, we grow nanotips in converging directions, eventually forming

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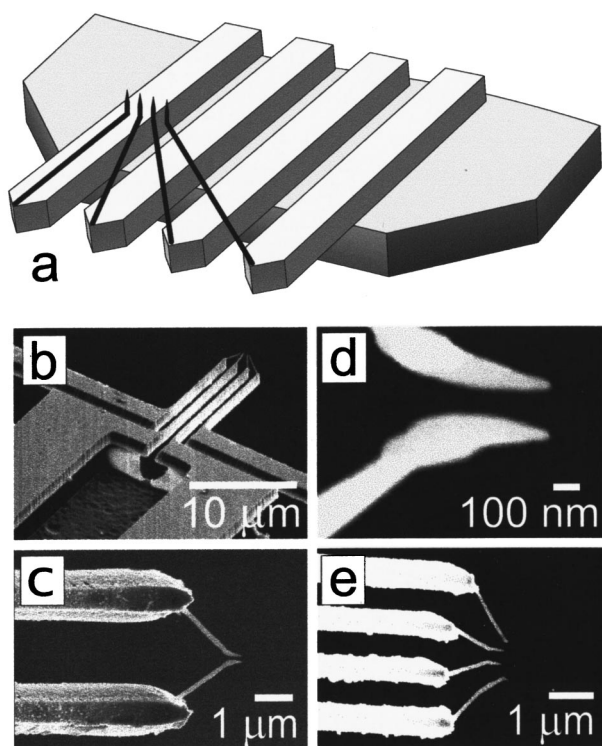


FIG. 1. (a) Schematic of the nanotips deposited in a plane perpendicular to the plane of the microcantilevers. (b) SEM image of microfour-point probe with $4\ \mu\text{m}$ spacing. (c) SEM image of nanotwo probe with $200\ \text{nm}$ spacing, imaged at an angle of 45° . (d) Close up of $200\ \text{nm}$ gap two-probe. (e) SEM image of nanofour-point probe with an average spacing of $350\ \text{nm}$.

multipoint probes with nanoscale electrode gaps, in a plane perpendicular to the plane of the microcantilevers [see Fig. 1(a)]. In Fig. 1(c) a SEM image of a two-probe with a $200\ \text{nm}$ gap is shown. To fine tune the gap and lengths of the tips, parallel secondary tips are added to the converging primary tips. The tips are grown in an alternating, iterative fashion, using successively shorter deposition times. The final adjustments are done with deposition times of about $5\ \text{s}$, allowing the gap and tip lengths to be tuned to within $10\ \text{nm}$ [see Fig. 1(d)]. In Fig. 1(e) an example of a four-point probe with an average spacing of $330\ \text{nm}$ is shown. Larger four-point probes have been fabricated with tip lengths of up to $3.5\ \mu\text{m}$ and gaps of $500\ \text{nm}$.

III. EXPERIMENTAL TESTS

For testing the nano four-point probes we used a silicon substrate covered with a $1000\ \text{\AA}$ thin gold film, located on an xyz stage for positioning with $100\ \text{nm}$ accuracy. The probe is mounted at an angle of 30° with respect to the sample, in the focal point of a high-resolution video microscope. A laser beam is directed towards the microcantilever, so that four spots of reflected light are visible in the video microscope.⁴ When the cantilevers touch down onto the sample, a slight deflection leads to a measurable change in reflection intensity, allowing accurate control of the contact pressure. Due to their small diameter, the tips cannot themselves be resolved optically.

Figure 2 illustrated the measurement setup. An electronic switch in turn connects each electrode to an alternating

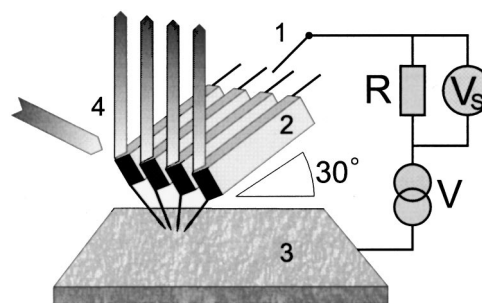


FIG. 2. Illustration of test setup. An electronic switch (1) in turn connects the electrodes to a two-point measurement circuit. The probe (2) is tilted 30° with respect to the sample (3). An xyz stage positions the sample with respect to the probe. A laser beam (4) reflected on the four microcantilevers allows monitoring of the deflection. The resistance is extracted from the voltage drop V_s over the known resistor R .

current voltage source V , which is connected to the sample through a series resistor R . When electrical contact is established to the sample, the voltage drop V_s across the series resistor can be used to determine the combined resistance, $R_s = (V/V_s - 1)R$, of the contact area, the electrode, the sample, and the electronic switch.

With this setup an upper limit for the individual resistance of the electrodes can be determined as a function of the vertical sample position (see Fig. 3). Upon contact [down arrow, Fig. 3(a)] the resistance drops from about $30\ \text{M}\Omega$ to $1\ \text{k}\Omega$ for electrodes 1, 2, and 3, which electrode 4 does not conduct. Upon retraction [up arrow, Figs. 3(a) and 3(b)] the release point is $100\text{--}300\ \text{nm}$ higher than the engage point. This hysteresis is due to sticking of the gold electrodes to the gold surface.

We test the mechanical strength of all nanoelectrodes by continually moving the sample towards the probe, until we provoke mechanical breaking of the tips, which is observed

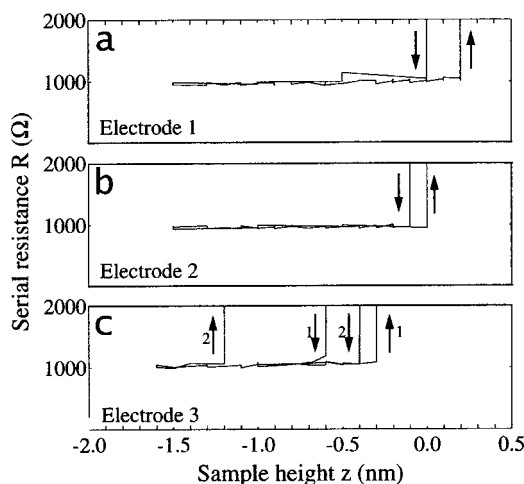


FIG. 3. Serial resistance as a function of sample height (arbitrary zero) for the three working electrodes. (a) The nanotip engages at $0.0\ \mu\text{m}$ but releases at $+0.2\ \mu\text{m}$, due to sticking. (b) Similar graph for electrode 2. (c) For electrode 3, two curves are shown. The first approach marked with 1 shows contact (down arrow) at $-0.6\ \mu\text{m}$ and release (up arrow) at $-0.3\ \mu\text{m}$. The second approach marked with 2 shows the contact occurring at $-0.4\ \mu\text{m}$ (down arrow), but here the probe clearly breaks at $-1.2\ \mu\text{m}$ (up arrow). For subsequent contacting approaches of electrode 3, an unstable contact is obtained at $-1.2\ \mu\text{m}$ rather than $-0.4\ \mu\text{m}$.

as a sudden increase of the resistance. As seen in Fig. 3(c), electrode 3 is engaged a second time (down arrow 2), and suddenly breaks during retraction (up arrow 2). In the subsequent measurements, the contact height for electrode 3 is displaced $0.8 \mu\text{m}$ further down than before the nanoprobe break. This critical deflection corresponds to a contact force of roughly 10^{-5} N. SEM images acquired afterwards confirm that the nanoelectrodes are broken off, while the supporting microelectrodes are still intact. This behavior is also observed for electrodes 1 and 2. Despite their small dimensions, the nanoprobes are thus sufficiently robust to sustain a contact force strong enough to deflect the cantilevers by more than $1 \mu\text{m}$. The 700 nm wide, 1500 nm tall microelectrodes typically break at a deflection of $3\text{--}4 \mu\text{m}$, which implies that the material composing the nanotips is substantially stronger than SiO_2 .

IV. SUMMARY

In summary, we show that it is possible to make multi-point probes with electrode spacings down 200 nm , using a combination of conventional microlithography and electron beam deposition. The new probe enables positioning of the source, drain, and voltage electrodes on the same micro-sized object, and thereby extends the four-point measuring technique to high spatial resolution scanning measurements. The use of flexible microcantilevers as base for the nanotips ensures that the electrodes can adapt individually to height differences on the surface. We have tested one such prototype and succeeded in establishing reliable, reproducible contact to a conducting surface.

Applications for such nanoscale multiprobe array include conductivity measurements on single atomic terraces and conducting polymer films,⁴ as well as magnetic and su-

perconducting grains, domains and thin films. Such a probe can be used to create high resolution conductivity maps of thin films, due to reduction of the spatial range of geometric effects compared to previous experiments.⁴ Several fascinating prospects of a nanoscale two probe for studying mesoscopic transport on surfaces and wires have been reported in literature.^{11,12} The small scale of the probe, however, requires high-resolution microscopy for accurate positioning on small objects, which can be accomplished using an *in situ* SEM.

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