Microfour-point probe for studying electronic transport through surface states

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Microfour-point probes integrated on silicon chips have been fabricated with probe spacings in the range 4–60 \( \mu \)m. They provide a simple robust device for electrical transport measurements at surfaces, bridging the gap between conventional macroscopic four-point probes and scanning tunneling microscopy. Measurements on Si(111) surfaces in ultrahigh vacuum reveal that the Si(111)-\( \sqrt{3}\times\sqrt{3}\)-Ag structure induced by a monolayer of Ag atoms has a four-point resistance two orders of magnitude lower than that of the Si(111)-7\times7 clean surface. We attribute this remarkable difference to direct transport through surface states, which is not observed on the macroscopic scale, presumably due to scattering at atomic steps. © 2000 American Institute of Physics.

The samples were 20\( \times \)3 mm\(^2\) n-type Si(111) with a nominal resistivity of 10–100 \( \Omega \)cm. The sample surfaces were patterned to generate large terraces during flashing. The patterning was done using a laser etching facility. To make large step-free terraces, grids of micron-sized small holes were etched with the laser, similar to the patterns used by Ogino. Grid spacings of 5, 10, 15, and 20 \( \mu \)m were used. The samples were heated resistively at 1250 °C in intervals of 10–60 s, while the chamber pressure was kept below 5 \( \times 10^{-9} \) Torr. The flashing procedure created step bunching due to electromigration. After a total of 3000 s of flashing, the step bunches were roughly aligned to the position of the original hole grid.

The two surface reconstructions investigated in this study were the Si(111)-7\( \times \)7 and the Si(111)-\( \sqrt{3}\times\sqrt{3}\)-Ag. Both have a surface state at the Fermi energy, and should thus be conducting. However, the dispersion of the half-filled state on the Si(111)-7\( \times \)7 surface is very low, and recent STM measurements indicate that this surface has a low conductance. The Si(111)-\( \sqrt{3}\times\sqrt{3}\)-Ag surface, on the other hand, has a band with strong dispersion at the Fermi level.

Macroscopic four-point measurements showed a 10% difference in conductance between the two reconstructions.

The position of single atomic steps was revealed by deposition of around 0.1 monolayers of Ag, which decorates step edges. It was thus confirmed that atomically flat terraces were created between the step bunches. The microscopic four-point probe was then brought into contact with the sample, as shown in Fig. 1(b). The total probe width is larger than the width of the terrace, so the electrodes are positioned on neighboring terraces.

Measurements on the Si(111)-7\( \times \)7 and Si(111)-\( \sqrt{3}\times\sqrt{3}\)-Ag surfaces are shown in Fig. 2. These measurements were performed with a probe with 20 \( \mu \)m electrode spacing in a region with 15 \( \mu \)m spaced step bunches. The four-point probe resistance was 24 k\( \Omega \) on the 7\( \times \)7 surface and 280 \( \Omega \) on the \( \sqrt{3}\times\sqrt{3} \) surface. The measured values were reproduced.

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We suggest two possible explanations. One is that a larger fraction of the current runs in the space charge layer for the microscopic probe and thus the effect of the space charge layer is more significant in the microscopic case. This is mainly a geometrical effect. The other possible explanation is that a larger fraction of the current runs in the surface states.

Concerning the space charge layer, we note that the clean Si(111) surface has the Fermi level pinned at 0.65 eV above the valence-band maximum.\textsuperscript{12} For an \( n \)-type 100 \( \Omega \) cm silicon crystal, the bulk Fermi level is approximately 0.75 eV above the valence-band maximum.\textsuperscript{13} The band bending beneath the 7\( \times \)7 surface is thus only about 0.1 eV, and so the space charge layer region has a conductance close to that of the bulk. Under these conditions, the sample can be approximated as a semi-infinite system with a uniform homogenous conductance. In this limit, the bulk resistivity \( \rho \) can be calculated from the four-point probe resistance \( R \) as \( \rho = 2 \pi s R \), where \( s \) is the electrode spacing of the probe.\textsuperscript{14} A resistivity value of 295 \( \Omega \) cm is deduced, which is comparable with the nominal bulk resistivity of the sample, and hence consistent with the semi-infinite approximation.

Since the four-point probe resistance falls two orders of magnitude after deposition of a monolayer of Ag, we can conclude that the surface channel effectively shorts out the bulk channel in this case. The Fermi level for the \( \sqrt{3} \times \sqrt{3} \) surface is pinned at a position 0.1 eV above the valence-band maximum,\textsuperscript{2} and thus the band bending is in this case 0.65 eV. The electrode spacing is considerably larger than the space charge layer thickness, as estimated from the nominal bulk resistivity. Therefore, we can approximate the space charge layer as an infinite two-dimensional sheet, and the surface sheet resistivity \( R_s \) can then be extracted for the \( \sqrt{3} \times \sqrt{3} \) surface as \( R_s = R/\ln 2 \). This yields a sheet resistance of \( 1.2 \times 10^3 \) \( \Omega \), or a corresponding conductance of \( 8.0 \times 10^{-4} \) \( \Omega^{-1} \). For comparison, macroscopic four-point measurements by Hasegawa \textit{et al.} show a difference in conductance between the \( \sqrt{3} \times \sqrt{3} \) and the 7\( \times \)7 surfaces of about \( 1.2 \times 10^{-4} \) \( \Omega^{-1} \).\textsuperscript{2} In Fig. 3, the microscopic and macroscopic results are displayed together with a curve depicting the theoretical variation of the space charge layer conductance \( \sigma_{sc} \) as a function of band bending, calculated from the bulk carrier mobilities \( \mu_n \), \( \mu_p \), and the excess carrier densities \( \Delta N \), \( \Delta P \) in the space charge layer.\textsuperscript{15}

\[
\sigma_{sc} = e \mu_n \Delta N + e \mu_p \Delta P.
\]

The surface conductance of the \( \sqrt{3} \times \sqrt{3} \) surface as measured with the microscopic four-point probe is nearly one order of magnitude higher than expected from band bending alone. We therefore conclude that the extra conductance measured with the microfour-point probe on the \( \sqrt{3} \times \sqrt{3} \) surface is due to conduction directly through the surface states.

We suggest two possible explanations for obtaining this result on the microscopic scale. One is that the scattering in the surface state channel is reduced due to the smaller number of atomic steps between the voltage probes. The other explanation is improved electrical contact to the surface states using the flexible microcantilever electrodes. Tests show that the electrodes leave only minimal damage over a contact area of \( -100 \times 100 \) nm\(^2\).
An estimate of the conductivity of the surface states can be deduced from the difference in conductance between the microprobe measurement and theoretical prediction for band bending alone, $7.6 \times 10^{-4}$ $\Omega^{-1}$, and the typical thickness of a reconstructed surface, 1 nm. This yields a resistivity of $1.3 \times 10^{-4}$ $\Omega$ cm, comparable with the resistivities of bulk metals such as bismuth. The $\sqrt{3} \times \sqrt{3}$ structure has no relation to metallic silver. Therefore, we emphasize that this metallic conductivity is not due to a thin metallic film, but is an intrinsic property of a surface reconstruction.

A more detailed theoretical analysis of this result will require modeling four-point probe measurements in realistic systems (not semi-infinite or planar), and including effects of step structure and finite contact area. In addition, comparison with theory will require a complete mapping of the band structures of these surfaces.

The technique presented here can clearly be extended to study other surface reconstructions, each having potentially unique electronic transport properties. The microfour-point probe is thus a useful addition to local probe techniques for applications in surface science and technology.

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