Lawrence Berkeley National Laboratory

Recent Work

Title

Quantum Mechanical Capacitance in a Scanning Tunneling Microscope

Permalink

https://escholarship.org/uc/item/0tj0p6k7

Authors

Botkin, D. Weiss, S. Ogletree, D.F. <u>et al.</u>

Publication Date 1995-02-01

Does Not

.BL-36845

Сору

Lawrence Berkeley Laboratory

Materials Sciences Division

To be presented at Ultrafast Electronics and Optoelectronics, Dana Point, CA, March 13, 1995, and to be published in the Proceedings

Quantum Mechanical Capacitance in a Scanning Tunneling Microscope

D. Botkin, S. Weiss, D.F. Ogletree, M. Salmeron, and D.S. Chemla

February 1995

U. C. Lawrence Berkeley Laboratory Líbrary, Berkeley

FOR REFERENCE

Not to be taken from this room



Prepared for the U.S. Department of Energy under Contract Number DE-AC03-76SF00098

DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California.

LBL 36845 UC-400/000

QUANTUM MECHANICAL CAPACITANCE IN A SCANNING TUNNELING MICROSCOPE

D. Botkin, S. Weiss, D. F. Ogletree, M. Salmeron and D. S. Chemla

Department of Physics University of California Berkeley, CA 94720

and

Materials Sciences Division Lawrence Berkeley Laboratory University of California Berkeley, CA 94720

February 1995

This work was supported by the Laboratory Directed Research and Development Program of Lawrence Berkeley Laboratory under the U.S. Department of Energy, Contract No. DE-AC03-76SF00098, and by ONR/DARPA under Contract No. N000-14.93.105.36.

Quantum Mechanical Capacitance in a Scanning Tunneling Microscope

D. Botkin, S. Weiss, D.F. Ogletree, M. Salmeron and D.S. Chemla

Materials Sciences Division, MS 2-300 Lawrence Berkeley Laboratory and Department of Physics University of California at Berkeley

Berkeley, CA 94720

Tel: (510) 486 - 5265 FAX: (510) 486 - 5530 e-mail: dbotkin@ux5.lbl.gov

<u>Abstract</u>

The nature of the ultrafast response of the tunnel junction in a scanning tunneling microscope is discussed.

Not

Quantum Mechanical Capacitance in a Scanning Tunneling Microscope

D. Botkin¹ S. Weiss², D.F. Ogletree² M. Salmeron² and D.S. Chemla^{1,2}

¹Department of Physics, University of California at Berkeley, Berkeley, CA 94720 ²Materials Sciences Division, MS 2-300, Lawrence Berkeley Laboratory, Berkeley, CA 94720

One of the most spectacular manifestations of quantum mechanics is tunneling between closely spaced conductors. The DC I-V characteristics in different types of tunnel junctions, such as metal-vacuum-metal junctions (MVM), metal-insulator-metal junctions (MIM), Josephson junctions, and quantum point contacts in a two dimensional electron gas, are well understood. High frequency transport in these systems is much less understood, and has been the subject of intense research in recent years[1,2].

The recently developed ultrafast scanning tunneling microscope (USTM) [3] is an ideal tool for the study of high frequency transport in the MVM or MIM system in a cross-sectional area on the order of 1 nm². In the USTM, an ultrafast photoconductive switch integrated with a scanning tunneling microscope (STM) tip assembly provides an ultrafast gate which temporally resolves the tunneling current. A "pump" laser pulse generates the surface excitation to be studied, and a "probe" laser pulse opens the sampling gate. An intriguing application of the USTM is the creation of movies of surface dynamics. At a fixed time delay between laser pulses, the tip can scan the surface, creating a snapshot of an ultrafast process. Such snapshots, viewed in succession, form a movie of surface dynamics. Before one can attempt to form movies, however, the USTM's response at a single point must be understood.

In what follows, we present experimental measurements of the high frequency response of the tunneling junction of a STM operating in air at room temperature. The tunnel junction is formed between the tip of the STM and the metallic surface of one strip of a coplanar transmission line on the sample. The STM tip position is fixed above a single point on the sample. A laser generates a picosecond voltage pulse on the transmission line, and the resulting tunnel current is monitored in time. The impedance of the tunneling gap can be deduced from the lineshape of the time-resolved tunnel current.

A typical signal measured by the USTM consists of a small AC component which varies with time delay and rides on a large DC background. The background is simply the DC tunneling current used for feedback, determined by the DC bias voltage and gap resistance. Fig 1a shows a 4.5 ps wide tunneling response to a 1 ps excitation pulse on the transmission line. The 10% to 90% rise time is 2 ps. The signal can also be measured while the tip is in ohmic contact, or crashed, with the sample. A typical crashed trace is shown in Fig. 1b. The time-resolved signal appears qualitatively different from the ohmic trace. In fact, the time-resolved signal's close resemblence to the derivative of the ohmic signal, shown in Fig. 1c., suggests that the tunnel junction has a partially capacitive response are very different from that expected for the geometrical capacitance of the junction.

1

To better quantify the tunneling capacitance, we model the tunnel junction as a resistor, R_i , and a capacitor, C_i , in parallel. With the approximations: (1) that the RC discharge time of the circuit is smaller than the voltage pulsewidth, and (2). that the temporal width of the on cycle of the photoconductive switch is smaller than the voltage pulsewidth, we find that the time resolved signal is given by:

$$S(\tau) \propto S_{cr}(\tau) + T_t \cdot S_{cr}(\tau) \qquad (1)$$

where $T_t = R_tC_t$ and $S_{cr}(\tau)$ is the crashed response. Eq. (1) states that the time resolved tunnel current can be fitted to $S_{cr}(\tau)$ and its derivative in time delay $S_{cr}'(\tau)$ with a single fitting parameter $T_t = R_tC_t$. This parameter has the dimensions of time delay, and is defined as the tunnel junction RC time constant. Since the tunneling gap resistance is known from DC measurements, we can extract the gap capacitance from the fitting parameter.

Fig. 2 is an example of such a fit. The solid line is the data of Fig. 1a. The fit (gray line) is



Fig. 1. Time resolved current crosscorrelation detected on the tip: (a) in tunneling range (5nA and 80mVsettings), and (b) when the tip is crashed into the sample. (c) is the time derivative of (b).

obtained, according to Eq. (1), from the crashed signal data of Fig. 1b and its derivative in τ . The tunneling capacitance extracted from the fit in Fig. 2 is $C_t = 6.7 \cdot 10^{-19}$ F. In a variety of measurements, we find C_t s varying from 10⁻¹⁹ to 10⁻²¹ F. In contrast, the geometrical tip-sample capacitance is estimated to be larger than 10⁻¹⁵ F, even when the radius of curvature of the tip is as small as 1000 Å. The geometrical capacitance

of the tip structure measured by conventional means at 20 kHz is 10^{-13} F. The dependence of C, on tip-sample

separation offers additional insight into the origin of this capacitance. This dependence is measured by retracting the tip and recording the time resolved tunnel current for different tip heights. In Fig. 3, we plot the extracted capacitance for three sets of measurements, each set taken with different combinations of tip and sample metallization or different excitation geometry, as a function of the tunneling gap conductance, G_t . We would prefer to plot C_t as a function of the tip-sample separation, or barrier width, d.



Fig. 2: Fit (solid gray line) to the time resolved data (solid black line) with the ohmic contact signal

However, in ambient conditions, the apparent inverse decay length and apparent barrier width is very different from a clean MVM junction, while the true inverse decay length is still represented by G_t . We find that the tunnel gap time constant, $\mathbf{RC} = \mathbf{CG}^{-1}$, depends only weakly on the barrier width. Since $G_t = G_0 e^{-\alpha(d-d_0)}$, where α is the inverse decay length, and G_0 is the conductance at a reference barrier width d_0 [4], the measured capacitance must vary as $C_t \propto e^{-\alpha(d-d_0)}$; in contrast, the geometrical capacitance can decrease no more rapidly than 1/d.

We conclude from the above discussion that (1) the time resolved tunnel current has a large capacitive component which originates from the

tunneling itself, (2) the extracted capacitance is very small and is not the geometrical capacitance of the junction and (3) the capacitance varies exponentially with barrier width. Therefore, the capacitance has a quantum mechanical origin.

Experimental evidence of quantum reactance in the quasiparticle current in Josephson junctions [1] and in quantum point contacts in two dimensional electron gas systems [2] has been demonstrated only very recently. In both cases, the reactance is explained by the theory of photon assisted tunneling (PAT) [5] as originating from virtual transitions between the two electrodes of the junction. According to the PAT theory, the reactance can be derived from the DC I-V relation by a Kramers-Kroning (K-K) transformation. Since the DC response varies exponentially with tip height. K-K а transformation will preserve this exponential dependence, in accordance with experimental



Fig. 3 Junction capacitance vs. conductance. Data represented by circles taken with brass tip on Au sample. Data represented by diamonds or triangles taken with Pt-Rh tip on Al sample.

observations. Preliminary calculations indicate that the magnitude of the PAT capacitance is comparable to our results. An additional contribution to the quantum reactance can arise from the "electrochemical capacitance" [6] which originates from incomplete screening of the electric field in the metal.

This work was supported by a Laboratory Directed Research and Development grant from the Lawrence Berkeley Laboratory, under the US Department of Energy, contract DE-AC03-76SF00098 and by ONR/DARPA under contract #N000-14.93.105.36

References:

- Q. Hu, C.A. Mears, P.L. Richards and F.L. Floyd. Phys. Rev. Lett. 64, 2945 (1990) and A.H. Worsham, N.G. Ugras, D. Winkler, D.E. Prober, N.R. Erickson and P.F. Goldsmith, Phys. Rev. Lett. 67, 3034 (1991).
- [2] L.P. Kouwenhoven, S. Jauhar, J. Orenstein, P.L. McEuen, Y. Nagamune, J. Motohisa and H. Sakaki, to appear in Phys. Rev. Lett.
- [3] S. Weiss, D.F. Ogletree, D. Botkin, M. Salmeron and D.S. Chemla, Appl. Phys. Lett. 63, 2567 (1993).
- [4] J.G. Simmons, J. Appl. Phys. 34, 1793 (1963).
- [5] R. Tucker, IEEE J. Quantum Electron. QE-15, 1234 (1979).
- [6] M. Büttiker, J. Phys.: Condens. Matter 5, 9361 (1993).

LAWRENCE BERKELEY LABORATORY UNIVERSITY OF CALIFORNIA TECHNICAL INFORMATION DEPARTMENT BERKELEY, CALIFORNIA 94720