A Highly Stable Electrochemical Scanning Tunneling Microscope

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We demonstrate a home-built electrochemical scanning tunneling microscope (ECSTM). The ECSTM exhibits highly stable performance. The drifting rates in \(XY\) and \(Z\) directions of the ECSTM are about 67 and 55.6 pm/min, respectively. Moreover, a specially designed scanner unit successfully solves the well-known problem of large leakage current in high humidity atmosphere. The mechanical structure of the ECSTM is described in detail. The excellent performances of the system are demonstrated by the measured STM images (in copper sulfate solution), including clean and well-ordered large area morphology of Au(111) and the atomically resolved image of highly oriented pyrolytic graphite.

Key words: Scanning tunneling microscope, Electrochemistry, Highly stable

I. INTRODUCTION

Since scanning tunneling microscopy (STM) was invented in the early 1980s [1], it has been presented to be a powerful technique for real space imaging of surface structures at the atomic scale. Excitingly, STM has been brought to be operated not only in ultra-high vacuum (UHV), but also in air and liquids [2–5]. However, the stability of STM is still a key problem even after such a long time development.

In the normal STM, the imaging piezoelectric scanner or the sample is directly fixed on the coarse approach motor rigidly [6]. The instability of the motor caused by its own and external disturbance will be transferred into the tunneling junction. Consequently, these will downgrade the stability of the STM, especially for the electrochemical STM, which needs to be equipped with a cell. Hence, the electrochemical scanning tunneling microscope (ECSTM) cannot be built as compact as a normal STM. So the mechanical drift of the ECSTM is larger than that of the normal STM. The instability of the ECSTM would not only affect the measurement of spectroscopy in \(Z\) direction, but also affect the time of measurement on a specific target and distort the imaged lattice structure in highly resolved imaging [7].

Besides, during the measurement of ECSTM, the evaporation of the solution will change the concentration of the solution. And the external contamination would gradually diffuse into the solution, resulting in the sample being polluted. These will lead to the experiment termination. However, if the ECSTM was protected in a sealed container, the evaporation of the solution would increase humidity of the container, resulting in the reduction of the insulation resistance between the tip and scanner electrode [8]. As a result, a variation of the scan voltages produced a proportional leakage current, which would seriously affect the quality of the imaging. How to solve this contradiction is a big challenge.

In this work, we present how we address these issues by adopting a stack piezoelectric motor to drive a detachable scanner unit for the coarse approach. The specially designed scanner unit successfully solves the well-known problem of large leakage current in high humidity atmosphere. The excellent performances of the ECSTM are demonstrated by the measured images: (i) clean and well-ordered large area morphology of Au(111) and (ii) the atomically resolved image of highly oriented pyrolytic graphite (HOPG) in spiral scan mode [9], which was the first spiral scan image in electrochemical environment.

II. STM DESIGN AND WORKING PRINCIPLE

Figure 1(a) is the schematic diagram of the ECSTM. The coarse approach motor (M) of the ECSTM is a piezoelectric stack motor. The structure of the motor will be described in detail in our later work. Its work principle is similar to that of the GeckoDrive [10]. However, the output force of the motor can reach \(\sim 8\) times that of the GeckoDrive. So it can carry more load.

The titanium frame (B) was machined by a piece of titanium plate. The scanner unit (Fig.1(c)) was installed underneath the stack motor. The scanner (S) was a
four-quadrant piezo tube with 4.35 mm outer diameter, 8.5 mm length and 0.5 mm wall thickness (EBL #3 material from Staveley). The scanner was fixed on a base (P) and then they were glued inside of a polished sapphire tube (E1) of 15.5 mm in length×7.5 mm OD×0.75 mm. There were five side wall slots of the base (P) to accommodate electrode wires of the scanner. The tunnel current wire went from the hole in the center of the base (P). The sapphire tip holder (Q) was glued on the free end of the scanner. A small stainless steel tube with 0.55 mm outer diameter and 0.28 mm inner diameter was mounted at the center of the tip holder, into which the tip was inserted. A sapphire gap (E2) with a properly sized hole in the center was fixed on E1 by epoxy resin. The top of the sapphire tube (E1) was capped with a 3.0 mm diameter polished sapphire ball. Scanner electrode wires and tunneling current wire were elicited from the six side wall slots on the top of the sapphire tube (E1). The sapphire tube was then sandwiched by a tungsten rail (C2) and a titanium rail cover (C1), in which the sapphire tube was pressed against the knife edges of the rail by a spring strip (R) between the sapphire tube and the rail cover.

Underneath the scanner unit was a quite large space (Fig.1(a)), in which the cell (F) was mounted. After the motor was sealed by a stainless steel sleeve (A), the ECSTM was installed into a trousers-like stainless steel circuit box (G). All the wires connected to the controller through its two legs. This would guarantee the scanner, motor and wires sealing, which could avoid the leakage current problem caused by high humidity above the solution.

The scanning part can be pushed (through the sapphire ball (D)) to implement the coarse approach by the stack motor. When the tested tunneling current reached the pre-set value, the coarse approach was finished. Then the motor can withdraw a proper distance from the scanning part, which prevented the instability of the motor due to thermal drift, external vibration, etc., from being transferred into the scanning part, resulting in a highly stable scanning part owing to its small size and excellent clamping structure. The scanning part can be withdrawn through a segment of soft rope between the scanning part and the motor. The sample stage was fixed on the steps of the frame via a string strip underneath. This minimized the mechanic loop of tunneling junction.

III. PERFORMANCE

In order to check how this ECSTM performs, we took a series of experiments. All the electrolyte used in this work was 50 mmol/L H2SO4 (from Merck, Suprapur grade) plus 1 mmol/L CuSO4, which were prepared using milli-Q water. All the tips were hand cut from 0.25 mm thick 80:20 Pt/Ir wires, which were coated with polymethylstyrene. Platinum and copper wires of 0.5 mm thick were used as the counter electrode and quasi-reference electrode, respectively.

The analysis of drift in the XY plane was carried out by the collection of sequential images in the same area. The working electrode was Au(111) of bead type, prepared by the Clavilier method [16]. Figure 2 shows large area constant current scans of 300 nm×300 nm taken at a 60 min interval. It is noted that the marked feature A are hardly drifted, showing very high stability. The quantitatively measured XY plane drifting values as a function of time are plotted with linear fittings in Fig.3, where each data point is the drifting distance of A in the stack motor. When the tested tunneling current reached the pre-set value, the coarse approach was finished. Then the motor can withdraw a proper distance from the scanning part, which prevented the instability of the motor due to thermal drift, external vibration, etc., from being transferred into the scanning part, resulting in a highly stable scanning part owing to its small size and excellent clamping structure. The scanning part can be withdrawn through a segment of soft rope between the scanning part and the motor. The sample stage was fixed on the steps of the frame via a string strip underneath. This minimized the mechanic loop of tunneling junction.

FIG. 2 Repeated scans of the same 300 nm×300 nm Au(111) area in the solution of 50 mmol H2SO4+1 mmol CuSO4 with a 60 min time interval. (a) 0 min, (b) 60 min, and (c) 120 min.
ECSTM [7]. Measurement of drift in the Z-direction was quantified through long-term acquisition of the Z-signal in the tunneling condition. Figure 3 provides an example of the measured drift in the electrochemical environment and reveals a rate of 55.6 pm/min.

Figure 4(a) compares the effect of atmospheric humidity on leakage current for a common and our home-built ECSTM. During the process of measurement, the common and our home-built ECSTM were protected in a sealed container. The humidity of the container nearly reached saturation with the evaporation of the solution. Black and blue curves exhibit the relationship between the scanning voltage and leakage current of the common and the home-built ECSTM in the dry atmosphere, respectively. For comparison, the red and green curves show the relationship between the scanning voltage and leakage current of the common and the home-built ECSTM in the high humidity atmosphere, respectively. For comparison, the red and green curves show the relationship between the scanning voltage and leakage current of the common and the home-built ECSTM in the high humidity atmosphere, respectively. For comparison, the red and green curves show the relationship between the scanning voltage and leakage current of the common and the home-built ECSTM in the dry atmosphere. It can be seen that in the high humidity atmosphere the leakage current of the home-built ECSTM is one order lower than that of the common ECSTM at the same scanning voltage (using 200 MΩ feedback resistance). The effect of such lower leakage current of the home-built ECSTM on the imaging can be ignored. So the specially designed scanner unit successfully solved the problem of large leakage current in high humidity atmosphere [8]. As the final test, high-resolution image of HOPG sample (as shown in Fig.4(b)) was obtained in spiral scan mode which can reveal the details of the drifting [9]. It was the first spiral scan image of ECSTM. The figure was scanned at 250 s/image in constant height mode. The scan size was 7.5 nm × 7.5 nm. The tip potential was at 204 mV and the working electrode was at 120 mV (reference to Cu/Cu2+).

IV. CONCLUSION

We present a home-built ECSTM with high stability. The XY plane and Z direction drifting rates are about 67 and 55.6 pm/min, respectively. A specially designed scanner unit could guarantee the high stability and greatly reduce the leakage by preventing the scanner and electrode wires exposed to the high humidity above the electrolyte. The excellent performances of the system are demonstrated by the measured ECSTM images of large area morphology of Au(111) and the atomically resolved image of HOPG, which is the first spiral scan image in electrochemical environment.

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