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# Scaling behavior of (0 0 1) and (1 1 1) Cu surfaces

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#### Abstract

Single crystalline Cu films are extensively used as buffer layers for a variety of applications. Cu films can be grown epitaxially on hydrogen-terminated Si(0 0 1) and (7  $\times$  7) reconstructed Si(1 1 1) substrates but the ultimate surface of the "as-grown" films tends to be quite rough. For most applications this constitutes a technical drawback. Using correlated reflection high-energy electron diffraction (RHEED) and STM data, we have found a dramatic smoothing of epitaxial Cu(0 0 1) surfaces by annealing the as-grown films in the 120–160 °C temperature range and somewhat less so for the Cu(1 1 1) films. Scaling analysis of the subsequent Cu growth on the annealed smooth surfaces yields a coarsening exponent of one-fourth for the (0 0 1) oriented films while this exponent is one-third for the (1 1 1) films, providing experimental data for a comparison of the same system film–substrate in these two orientations. © 2002 Elsevier Science B.V. All rights reserved.

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# 1. Introduction

Buffer layers are often used to stabilize the growth of epitaxial films. In particular, Cu buffer layers grown on silicon substrates are used extensively for subsequent growth of epitaxial magnetic multilayers. Unfortunately the appearance of roughness as a result of noise, strain and step-edge barriers is an important challenge to the development of applications of epitaxial metal layers. In the case of heteroepitaxy, strain due to lattice mismatch and interfacial chemical reactions can also play an important role in roughening [1].

Interestingly, several authors indicate that copper can be grown at room temperature as a single-domain

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film on  $(0\ 0\ 1)$  Si–H or  $(1\ 1\ 1)$  Si-7 × 7 surfaces, while it is not possible to grow single-domain films on  $(1\ 1\ 1)$  Si–H or reconstructed  $(0\ 0\ 1)$  Si surfaces [2,3]. We describe later our results of an in situ study comparing the mass transport kinetics of singledomain  $(0\ 0\ 1)$  Cu films epitaxially grown on hydrogen-terminated Si $(1\ 0\ 0)$  and Cu  $(1\ 1\ 1)$  films epitaxially grown on 7 × 7 reconstructed Si $(1\ 1\ 1)$ , as well as the scaling behavior of growth parameters such as roughness and coarsening of the surface features.

In order to achieve smoother surfaces it is customary to anneal in situ the "as-grown" films. The annealing temperature must be high enough to provide enough energy to overcome step-edge barriers. However, in the case of Cu–Si, the interaction is relatively strong due to p–d hybridization in the Cu–Si intermixing region that results in silicide formation, even at 100 K [4]. The thickness of this silicide is strongly temperature dependent, thus limiting the annealing

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temperature range as well as the minimum thickness of the film to be annealed. We will show here that it is possible to obtain smooth single-domain copper films in both (0 0 1) and (1 1 1) orientations, grown on Si, via a mild annealing in a temperature range below predominant silicide formation. We have found that films of thickness ~100 nm or thicker do not exhibit silicide formation at the surface after the annealing process. A study to determine the lower limit of the film thickness capable of sustaining the annealing process without compromising the surface quality is under way.

# 2. Experimental

Our studies were performed in a molecular beam epitaxy (MBE) chamber equipped with an in situ ultrahigh vacuum (UHV) scanning tunneling microscope and a real-time reflection high-energy electron diffraction (RT-RHEED) imaging system [5]. Copper was evaporated from a K-cell at a rate of 0.2 Å/s. The films were grown under UHV conditions.

The  $(1\ 0\ 0)$  copper films were grown at room temperature (RT) on Si(0 0 1) substrates cleaned ultrasonically with acetone and alcohol for 15 min each and etched with HF (1 or 10% v/v) for 30 s. The surface of the substrates so treated is known to be hydrogen terminated. [6] Such surfaces are known to be inert for several minutes in air and for several hours in UHV at RT. This surface passivation is believed to be due to hydrogen termination of the dangling Si bonds, which renders the surface chemically stable.

The  $(1\ 1\ 1)$  copper films were grown at RT on  $(7 \times 7)$  reconstructed  $(1\ 1\ 1)$  Si. The Si $(1\ 1\ 1)$  substrates were prepared with the low temperature cleaning method proposed by Ishizaka and Shiraki [7]. This process consists of two steps: (a) wet chemical treatment to etch the contaminated surface layers and to form a contamination-free passivative oxide film on the Si substrate, followed by heating under UHV conditions at temperatures below 900 °C to remove the oxide film through thermal etching and obtain an atomically clean surface. The RHEED pattern for Si substrates treated in this way typically shows the  $(1\ 1\ 1)\ 7 \times 7$  superstructure, indicating an atomically clean surface.

### 3. Discussion

# 3.1. The (0 0 1) Cu films

During the initial stages of growth, the RHEED pattern of silicon faded and, shortly thereafter, the crystalline structure of copper became apparent. The in-plane crystallographic axes of copper are rotated  $45^{\circ}$  with respect to the silicon lattice as has been reported previously and ex situ cross sectional TEM and XRD corroborated the single-domain quality of the (0 0 1) oriented films [6]. After ~2000 Å of Cu deposition, the RHEED pattern had a symmetrical spotty appearance characteristic of three-dimensional growth (inset (a) in Fig. 1).

The film was then annealed by increasing the temperature from room temperature to 443 K (where the copper RHEED pattern fades) at a rate of 1 K/min. The heating process showed its effect through a sharpening of the copper streaks, indicating that the crystalline order has improved significantly as a result of the annealing (inset (b) in Fig. 1). A plot of 1/full width at half maximum (FWHM) of the specular RHEED streak versus temperature, shows two distinct regimes: A gradual narrowing for T < 395 K, and a more rapid evolution for T > 395 K. (Fig. 1) We have also observed that for T > 445 K, the RHEED pattern becomes very diffuse indicating that the surface becomes rough again marking the onset of silicide formation. At approximately 470 K the copper. RHEED pattern completely disappears. Subsequent



Fig. 1. Plot of the inverse width (pixels) of the RHEED central streak as a function of temperature. Inset: RHEED images of a 200 nm thick Cu film, (a) as-grown and (b) after annealing at 423 K.



Fig. 2. (a) STM image of the (0 0 1) Cu film as-grown; (b) STM image of the same film after annealing. Both images are  $300 \text{ nm} \times 300 \text{ nm}$  in size.

films were annealed isothermally in the fast regime (T > 390 K but below the silicide onset temperature, T < 445 K). Fig. 2(a) shows the STM image after growth at RT. The rms roughness of the surface is  $\sim 70 \text{ Å}$  and the grain size is small ( $\sim 160 \text{ Å}$ ) due to interfacial strain (the lattice mismatch for an in-plane rotation of  $45^{\circ}$  between the Si and Cu lattices is  $\sim 6\%$ ) and limited diffusion kinetics that lead to deviations from ideal epitaxial growth. Fig. 2(b) on the other hand, corresponds to the STM image after annealing the film at 423 K for 4 h. The surface of the film now has large islands (average size 800 Å) and a mean rms roughness of  $\sim 10 \text{ Å}$ .

In order to analyze the smoothing kinetics of this system, we followed a similar analysis to that performed by Zuo et al. [8]. During the annealing process, the average terrace width on the film surface L(t) changes in time following a  $t^{1/3}$  law, consistent with kinetics driven by the line tension of curved step edges [9]. For T > 360 K the size of the initial terrace width  $L(t_0)$  is not negligible and the n = 1/3 growth law should read as follows:

$$L^{3}(t) - L^{3}(t_{0}) = A(T)(t - t_{0})$$
(1)

where  $A(T) \propto (1/T) \exp(-(E_{act})/k_{B}T)$ , is a temperature-dependent factor. The terrace size is inversely proportional to the width ( $\Delta$ ) of the RHEED specular streak.

Thus, we plotted our RHEED data as  $[1/\Delta(t)]^3 - [1/\Delta(t_0)]^3$  versus  $(t - t_0)$ . In such plots, the exponents for the time variable  $[(t - t_0)$  in Eq. (1)] extracted for all temperatures were in the range  $0.98 \pm 0.02$ , consistent with the n = 1/3 law. The pre-factor A(T) was obtained for all temperatures from a least-squares fit

to the data. An Arrhenius plot of  $\log[TA(T)]$  versus 1/T yielded an activation energy of  $0.40 \pm 0.04$  eV. The latest reported value for the activation energy for surface diffusion is  $0.36 \pm 0.03$  eV [10]. Therefore, our value for the activation energy indicates that the fast smoothing of the surface is dominated by activated diffusion over step barriers in the temperature interval considered.

In order to further characterize the nature of the annealed Cu/Si heteroepitaxial surface we studied the scaling behavior of the growth after further copper deposition. The rms roughness (interface width)  $\xi$  for length *L* and average thickness  $\langle h \rangle$ , for a system containing *N* sites with single valued heights  $h_i$ , is defined by the following expression [11]:

$$\xi(L,\langle h\rangle) = \left[N^{-1}\sum_{i}^{N}(h_{i}-\langle h\rangle)^{2}\right]^{1/2}$$
(2)

This theory predicts that when  $\langle h \rangle \rightarrow 0$ ,  $\xi$  scales as:

$$\xi(\langle h \rangle) \propto \langle h \rangle^{\beta} \tag{3}$$

where  $\beta$  is the dynamic roughness exponent and when  $\langle h \rangle \rightarrow \infty$ , it scales as:

$$\xi(L) \propto L^{\alpha} \tag{4}$$

where  $\alpha$  is the static roughness exponent. The exponent  $\beta$  was determined from a series of STM images obtained after the deposition of 3-15 monolayers (MLs) of Cu on the smooth surface obtained after annealing. The exponent  $\alpha$  on the other hand, was obtained from a single image obtained after 2000 Å of further Cu deposition. Accordingly,  $\alpha$  was derived from Eq. (4) where L corresponds to a segment of the STM line-scan measured along a direction parallel to the surface. The plot log  $\xi$  versus log L exhibited a linear region for values of L smaller than the island size  $(d_s)$  and after that it saturated. The slope of this straight line was found to be  $1.03 \pm 0.03$ . The exponent  $\beta$  was evaluated from Eq. (3). From a plot of log  $\beta$ versus log  $\langle h \rangle$ , a straight line is obtained with a slope  $\beta = 0.25 \pm 0.01.$ 

We also measured the length-scale associated with coarsening. As  $\langle h \rangle$  increases, the average island size  $d_s$  increases according to:

$$d_{\rm s} \propto \langle h \rangle^{\beta/\alpha} \tag{5}$$



Fig. 3. (a) RHEED image of the  $7 \times 7$  Si(1 1 1) surface; (b) STM image of the surface of a (1 1 1) Cu film epitaxially grown on the  $7 \times 7$  (1 1 1) Si surface; (c) RHEED image of the (1 1 1) film after annealing; (d) STM image of the surface of the annealed Cu film. The bar on the STM images corresponds to 50 nm.

From the plot log  $d_s$  versus log  $\langle h \rangle$  a value for  $\beta/\alpha = 0.25 \pm 0.01$  was obtained, and using the value obtained previously for  $\alpha \ (\approx 1)$  we estimate  $\beta = 0.25$ , consistent with the value derived from Eq. (3). Our values agree with those reported by Zuo et al. [8] for homoepitaxial growth of  $(0\ 0\ 1)$  copper at RT and also with theoretical models [12].

#### 3.2. The (1 1 1) Cu films

Fig. 3(a) shows the RHEED pattern of the  $7 \times 7$  reconstruction of the (1 1 1) Si surface prior the growth. After 10 ML of copper growth, the  $7 \times 7$  Si(1 1 1) pattern has disappeared, and only the fundamental reflections of Cu can be observed. Here, the epitaxy is controlled by the formation of an interfacial compound with a thickness of a few MLs. [3] The RHEED pattern shows only the fundamental reflections of the epitaxial Cu layers. This pattern has a spacing that is commensurate with the Cu lattice spacing and a unit cell that is not rotated with respect to the underlying Si substrate. After further deposition

the RHEED pattern shows longer and sharper streaks, as determined by their smaller FWHM, indicating smooth single crystal film growth. Fig. 3(b) shows the STM image of the surface as grown for a film 100 nm thick. We notice that the surface is mounded with no special features.

Annealing the films to improve the surface roughness was constrained even more than for the  $(0\ 0\ 1)$ case, as the film grew rougher probably due to increase of the silicide region at the interface during growth. In the temperature interval 298 K < T < 373 K, the activation energy measured for the smoothening was  $1.1 \pm 0.03$  eV. Despite this somewhat high activation energy for mass transport, a significant improvement of the surface roughness was observed (Fig. 3(c)). Fig. 3(d) shows an STM image of the surface where hexagonal terraces are now evident. Zuo et al. [8] also found an activation energy of 1.10 eV in the temperature range 333 K < T < 368 K for (0 0 1) Cu. The larger value measured for the activation energy for mass transport in H-free systems, suggests that the H termination on the (001) silicon/copper interface may speed up the kinetics at the  $(0\ 0\ 1)$  Cu surface, although at the present time the mechanism for this process is not understood. Scaling studies were carried out for subsequent growth of copper on the (1 1 1) annealed surface. These studies yielded a coarsening exponent of 0.333 in agreement with theoretical predictions for the (1 1 1) orientation [13].

#### 4. Conclusions

We have achieved smooth, epitaxial single-domain copper films MBE grown on Si substrates. The annealing temperature interval to smoothen the films is severely constrained due to silicide formation at the film–substrate interface. We have not observed silicide presence at the surface of these films, for film thickness ~100 nm and above. The activation energy measured for mass transport is considerably smaller for the (0 0 1) surface than for (1 1 1), suggesting that H termination on the (0 0 1) silicon/copper interface may intervene during the annealing and speed up the kinetics at the (0 0 1) Cu surface. Scaling studies yield coarsening exponents of one-fourth for the (0 0 1) surface and one-third for the (1 1 1) surface in agreement with theoretical models, and providing for the first time an experimental comparison of the growth for the same material and substrate.

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