Thermal-Hydrogen Promoted Selective Desorption and Enhanced Mobility of Adsorbed Radicals in Silicon Film Growth

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Car-Parrinello simulations and static density-functional theory calculations reveal how hydrogen promotes growth of epitaxial, ordered Si films in plasma-enhanced chemical vapor deposition at low-temperature conditions where the exposed Si(001)-(2 × 1) surface is fully hydrogenated. Thermal H atoms, indeed, are shown to selectively etch adsorbed silyl back to the gas phase or to form adsorbed species which can be easily incorporated into the crystal down to \( T \sim 200 \, ^\circ \text{C} \) and start diffusing around \( T \sim 300 \, ^\circ \text{C} \). Our results are well consistent with earlier experiments.

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Plasma-enhanced chemical vapor deposition (PECVD) is a very popular technique for growing silicon films. The possibility of reaching high growth rates at moderate temperatures has made PECVD particularly appealing for generating amorphous (a), microcrystalline (\( \mu \)c), and nanocrystalline (\( nc \)) films, of particular importance, e.g., in the production of solar cells [1]. By reducing the growth rate [2] or by raising the substrate temperature [3] also epitaxial silicon films of good quality can be obtained. This versatility surely makes PECVD of extreme scientific interest. Among the large amount of information coming out of decades of experiments and production, one phenomenon has captured particular attention, leading to several, sometimes controversial, interpretations. Increasing amounts of hydrogen impinging the surface were indeed shown to enhance crystallinity both during growth, and/or in post-growth treatments [2,4,5]. The observation of a simultaneous decrease in the growth rate, and/or of the thinning of the samples during hydrogen exposure, strongly suggested hydrogen to be able to preferentially etch noncrystalline regions [2,6]. A further, possibly complementary channel for H-induced crystallinity in post-growth exposure was also proposed [5]. While very many indications for possible competing mechanisms were given based on experimental observations, the complexity of the problem, in terms of geometry, chemistry, and the multitude of species involved, has prevented so far the development of a parameter-free quantitative modeling of the process at the atomic scale able to shed full light on the phenomenon.

In this Letter we shall focus on the role played by hydrogen during silicon-film growth starting from a fully hydrogenated Si(001)-(2 × 1) representative of the initial stages of low-temperature deposition, where thermal H desorption is not activated [7]. Our study has been motivated by the wide interest in hydrogen-induced crystallization, and, more specifically, by the seminal paper of Tsai et al. [2] in which it is demonstrated the possibility of growing epitaxial Si films with PECVD at a very low temperature \( T = 200 \, ^\circ \text{C} \), provided that silane is introduced in the reactor strongly diluted with \( \text{H}_2 \). In their paper, Tsai et al. suggested preferential etching of nonepitaxial adsorbed species to cause the improved quality of the grown sample. Hydrogen etching is proposed to be selective to such an extent as to remove any wrongly located adsorbate, while leaving already epitaxial species at the surface. Based on \textit{ab initio} calculations here we shall show that etching does indeed play a key role, but through a much richer variety of induced surface phenomena including promotion of epitaxial sites and surface diffusion. As demonstrated in Refs. [8,9], impingement of SiH\(_3\), the dominant radical in PECVD under several working conditions [10,11], easily produces a surface dangling bond on Si(001)-(2 × 1):H to which a second incoming silyl molecule can stick forming the stable, but not epitaxial, configuration reported in Fig. 1 (inset 2Ha). We have investigated possible paths out of this state using static density-functional theory calculations with ultrasoft pseudopotentials [12] as implemented in the plane-wave \textsc{vasp} code [13,14]. We adopted Perdew and Wang [15] gradient corrected exchange-correlation functional. Spin-polarization was taken into account. We used a slab of 6 layers with 16 atoms per layer and a vacuum region of ~10 Å. The surface was passivated by hydrogen (both at the top, in the dimerized configuration, and at the bottom). A \( 2 \times 2 \times 1 \) Monkhorst-Pack \( k \)-points mesh was used and the energy cutoff was set to 240 eV. In geometry optimizations all of the atoms were allowed to relax except for the two bottom Si layers that were kept frozen to bulk positions. Activation energies were computed using the nudged elastic band technique [16] with the climbing image refinement [17]. Convergence at local-minima (saddle points) was achieved when forces were less than 0.005 eV/Å (0.02 eV/Å). Dozens of barriers were sampled in order to understand the full kinetic scenario characterizing possible evolution out of the initial state. The result of these calculations was rather clear: no barrier out of state 2Ha below \( E_a \sim 2.2 \, \text{eV} \) seems to exist. Using harmonic tran-
After running 81 simulations with atomic H impinging the substrate with a thermal energy of 0.1 eV, we were able to build the cross section displayed in Fig. 1. Events can be divided into three main categories, which occur roughly with the same probability. Hydrogen can etch SiH$_3$ by an Eley-Rideal mechanism coming back to the gas phase as silane (region indicated with boxes in Fig. 1 with a cross section $\sigma \sim 3.8 \, \text{Å}^2$, the sampled surface cell being 20.3 Å$^2$), it can capture one hydrogen from the adsorbed silyl leaving an adsorbed SiH$_2$ in the configuration 1Hb displayed in Fig. 1, or, beside simple recoil, it can etch an adjacent surface H atom leaving a highly reactive dangling bond (for instance, configuration 1Hc in Fig. 1). These results show that etching is indeed possible with some non-negligible probability. Is it also strongly preferential as suggested in Ref. [2]? To answer this question, we have recomputed the etching cross section when SiH$_3$ is adsorbed in a weaker bound configuration. In particular, we have positioned the adsorbed silyl bound to a silicon atom of the second layer, in the configuration highlighted in Fig. 2. This configuration has been identified as a possible adsorption site from ab initio static calculation [20] and from MD simulations of the impact of SiH$_3$ radicals on Si(001)-(2 × 1) : H [9]. The cross section for H-induced etching is reported in Fig. 2: it strongly raises passing from $\sigma \sim 3.8 \, \text{Å}^2$ for the stable configuration (Fig. 1) to a value of $\sigma \sim 8.0 \, \text{Å}^2$, confirming the selectivity of the process as proposed in Ref. [2].

However, if selective etching explains how noncrystalline features can be removed, it is not sufficient to understand how an epitaxial film can actually grow. Indeed, we...
still need to explain how incorporation into epitaxial sites is possible. Let us consider the other two products (beside etched silane) of SiH₃ interaction with impinging H (configurations 1Hb and 1Hc of Fig. 1). After computing a set of possible barriers out of these states, the following scenario emerged. From configuration 1Hb, the system can incorporate into the crystal by directly jumping to the epitaxial configuration 1Ha surmounting a barrier of 0.64 eV. However, a lower-energy mechanism exists, leading to configuration 1Hc. Considering the relative energy of the three involved local minima (reported in the figure), the effective barrier for reaching the epitaxial site starting from 1Hb is of ~1 eV. Using again Eq. (1) for \( T \approx 200 \, ^\circ C \), one finds \( \tau \approx 4.6 \, ms \). At any deposition rate \( \Phi \ll 1/\tau \approx 200 \, ML/s \), it is then possible to grow epitaxial films. Considering the very slow growth rate used in Ref. [2], the reported high degree of crystallinity is fully consistent with our findings. The here presented incorporation process is very similar to the one proposed in Ref. [21] (silicon growth) and also in Ref. [22] (diamond growth). Notice, however, that here the whole set of processes leading to final incorporation from the initial radical adsorption site were considered and quantified.

While incorporation is the key step to grow epitaxial films, it is often also desirable to obtain films with small surface roughness. To this goal, fast diffusion of adsorbed species is needed [23,24]. While no detailed information about the roughness of the films grown in Ref. [2] is given, we find it useful and interesting to provide a theoretical investigation of surface mobility which could be exploited in future experiments aiming at controlling both epitaxy and surface corrugation. From site 1Hc, the silyl can diffuse as a whole along the dimer row to configuration 1Hc’. While the barrier for this event is of ~1.5 eV, actual long-range diffusion would imply visiting again the stablest site, yielding an effective \( E_a \simeq 1.64 \, eV \). This value corresponds to a diffusive time \( \tau_D \) of several hours at \( T = 200 \, ^\circ C \), mobility starting to become relevant at around 400 \, ^\circ C (\( \tau_D \simeq 0.1 \, s \) is here and below taken as an indicative threshold for defining the onset of diffusion, the estimate being relevant for deposition fluxes \( \leq 10 \, ML/s \)). However, one should take into account that while the adsorbate is diffusing under actual growth conditions, impinging H atoms could saturate the dangling bond which is present in any of the visited configurations. This would be detrimental since, as we have shown, no diffusive event takes place in the absence of the dangling bond. In this case (or, more in general, if the temperature is too low for activating even a single of the 1.64 eV jumps), we might ask what happens if a further H impinges on configuration 1Ha. Following the same procedure used to generate Fig. 1 the cross section illustrated in Fig. 3 is found. The region with extension \( \sigma \simeq 19.3 \, \AA^2 \) marked with upwards triangles indicates where incoming H removes one of the SiH₃ hydrogens or the H saturating the underlying dimer, leaving the substrate as H₂ and creating the configurations 0Ha or 0Hc (Fig. 3). The latter evolves into 0Hb almost immediately, since a barrier of only 0.05 eV must be surmounted. A quick comparison with Ref. [25] reveals 0Ha and 0Hb to be two of the local minima visited by a Si adatom when diffusing on Si(001)-(2 × 1) : H. This means that H-induced removal of 1 H effectively transforms the adsorbed SiH₃ in 1Hb into a simple adatom diffusing on a fully hydrogenated environment. As demonstrated in Ref. [25] using the local density approximation, such an adatom is very mobile, since an effective barrier \( E_D \simeq 1.2 \, eV \) characterizes diffusion along dimer rows. By repeating the calculation along the full path proposed in [25] with our gradient corrected exchange-correlation functional approach, we find only a slightly higher estimate \( E_D \simeq 1.4 \, eV \). In the present contest, two important conclusions can be made. First, thanks to the etching process, diffusion down to \( T \simeq 300 \, ^\circ C \) is allowed. Moreover, the whole diffusive process does not need additional freeing of dangling bonds in the neighborhood of the moving Si atom.
Therefore, surface diffusion can be sustained along extended paths also in the presence of continuously incoming H. Let us now analyze the remaining simulations. When H sticks in the left region of the sampled impact area of Fig. 3 configuration 2Hc (or more rarely 2Hd) is obtained, which can evolve with a barrier of \(\sim 0.5\) eV to the starting and motionless configuration 2Ha, ready to be further on etched. Alternatively, the epitaxial minimum 2Hb displayed in Fig. 3 is reached (when H sticks on the right region of the impact area and saturates the dangling bond). Here the adsorbed radical, while perfectly epitaxial, is totally trapped (see also Ref. [21]) so that it could be interesting to monitor its behavior under further H impingement. This does not require further simulations, since, due to the close resemblance with configuration 1Ha we can exploit the left portion of the cross section of Fig. 3. Three different outcomes are thus possible: one of the H atoms saturating the dangling bond of the same Si atom to which SiH\(_2\) is bound can be etched, restoring configuration 1Ha; alternatively, one of the 2 H atoms of SiH\(_2\) is etched, or, the incoming H atom is inserted into the Si-Si bond, similarly to configuration 2Hc of Fig. 3. While we have not completely explored possible further evolution, we believe none of the above possibilities to open up channels for faster mobility.

Hydrogen-induced enhanced mobility of ad-species was invoked in Ref. [26] to explain the experimentally observed strong decrease in film roughness with increasing H pressure. Although it is tempting to relate our theoretical estimation of the there observed slow decay of the etching rate with etching by atomic hydrogen is described in Ref. [28].

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Hydrogen-induced enhanced mobility of ad-species was invoked in Ref. [26] to explain the experimentally observed strong decrease in film roughness with increasing H pressure. Although it is tempting to relate our theoretical results also with this evidence, we note that in such work \(a\) and \(\mu c\) films were considered, so that the barriers here computed for the crystalline (001) surface should not be considered as more than indicative. A more direct confirmation of the predictive power of our calculations can be perhaps found in the etching experiments of Ref. [27], where both the here-presented H-induced mechanisms (silyl removal as silane, \(H_2\) removal leaving adsorbed SiH\(_2\)) were clearly identified. A further study of silyl etching by atomic hydrogen is described in Ref. [28].

The there observed slow decay of the etching rate with decreasing temperature seems fully compatible with our estimation of \(\sim 0.05\) eV, obtained performing a nudged elastic band calculation with the same procedure followed for computing the diffusion paths.

In summary, we have shown that hydrogen can promote ordered growth of fully hydrogenated PECVD silicon thin films through several channels, including etching-promoted radical incorporation into epitaxial sites (activated around \(T \sim 200^\circ\)C), etching-promoted diffusion \((T \approx 300^\circ\)C), and strongly selective etching of metastable adsorption sites. Our prediction for the onset of epitaxy, based on density functional theory, is consistent with experiments. Because of the complexity of the growth process a closer comparison with experiments, e.g., in terms of growth rates and/or surface roughness, still demands for further calculations and extensive modeling. We believe that the identification of elementary processes along with their activation energies and/or cross sections as presented here is of great value for the development of comprehensive kinetic Monte Carlo schemes, strongly needed to close the gap between atomic-scale and macroscopic measurements and gain a full understanding of PECVD growth.

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