Alloying mechanisms for epitaxial nanocrystals

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The different mechanisms involved in the alloying of epitaxial nanocrystals are reported in this letter. Intermixing during growth, surface diffusion and intra-island diffusion were investigated by varying the growth conditions and annealing environments during chemical vapor deposition. The relative importance of each mechanism was evaluated in determining a particular composition profile for dome-shaped Ge:Si (001) islands. For samples grown at a faster rate, intermixing during growth was reduced. Si surface diffusion dominates during H2 annealing whereas Ge surface diffusion and intra-island diffusion prevail during annealing in a PH3 environment.

In coherently-strained epitaxial islands, the most important factor that determines island size and stability is composition. Composition variations inside epitaxial islands will substantially influence their structural properties and, as a consequence, the electronic properties of epitaxial nanocrystals.[1]. The composition profile of SiGe islands has only recently been measured,[2,3,4] and its origin depends on kinetic and thermodynamic contributions, which sometimes are difficult to separate. In a careful and detailed study,[5] some of the pathways for Si and Ge intermixing have been investigated in Molecular Beam Epitaxy (MBE) grown islands. A series of experiments was carried out at different temperatures and with subsequent annealing steps in Ultra High Vacuum (UHV). The main result for the reported experimental conditions was that surface diffusion of Si and Ge was the dominant mechanism determining the island composition profile. It is known, however, that annealing and growth in different ambient conditions (i.e., UHV, as opposed to H2 or PH3 environments) selectively changes the surface mobility of adatoms.[6,7]. As a result, alloying in the presence of gases can proceed privileging selected mechanisms during Chemical Vapor Deposition (CVD) growth.

Bulk diffusion requires the formation of vacancies and/or interstitials.[8] For buried 2D SiGe layers, diffusion of Ge was found to increase with the Ge content and compressive stress.[9]. Although the amount of diffusion inferred from these results extrapolated to 600 °C is negligible for unstrained material, the activation energies depend strongly on strain. For the case of sub monolayer coverages, even at temperatures around 500 °C, Ge diffusion and intermixing into Si surfaces has been theoretically predicted and was observed by high resolution Rutherford Back Scattering (RBS) before the first monolayer of material was completely deposited.[10,11]. In self-assembled islands, a significant amount of strain is present and the fact that high index facets and edges (and hence defects) make up the surface provides a larger number of pathways for intermixing compared to a 2D film. Alloying in coherently-strained nanocrystals needs to be investigated in more detail to understand island formation and evolution. In order to comprehend the composition profiles, it is imperative to vary the kinetic and thermodynamic components individually in experiments to elucidate the mechanisms that lead to intermixing for a particular growth condition.

The primary mechanisms that modulate the composition profile of self-assembled islands are: a) exchange reactions between Si and Ge during island growth, defined by the attachment and detachment of atoms between the crystal and 2D adatom gas; b) surface diffusion of both Si and Ge adatoms; and c) diffusion of Ge and Si atoms within the islands (intra-island diffusion), excluding the surface adatoms. Inter-island diffusion, which is a special case of surface diffusion, is not covered here in detail, but is also a crucial component for the final island composition profile. Changing the growth and annealing conditions allows us to privilege one mechanism at a time, but not rigourously supress the other two. The goal of this work is to evaluate the relative importance of each mechanism by comparing samples grown by CVD and annealed under different conditions. For instance, comparing samples grown at different rates affects all mechanisms, but more effectively a and b. Annealing in a H2 environment decreases surface mobility of both Ge and Si compared to annealing in UHV,[6], yet does not completely stop the surface diffusion of either species. Thus it allows both mechanisms b and c to be investigated. Annealing in a PH3 environment substantially reduces Si surface diffusion, yet has little effect on Ge surface diffusion. This different behavior occurs because the P-Si bond is stable (bond enthalpy equal to 364 ± 7 kJ/mol) whereas the P-Ge bond is unstable.[12]. In this case, mechanism c dominates for both Ge and Si species, and mechanism b persists for Ge adatoms.

Four samples with nominally the same Ge deposition thickness of 12 eq-ML were grown at 600 °C in a H2 ambient in a commercial CVD reactor on 150 mm diameter Si (001) wafers. The conditions were chosen to produce dome-shaped-islands. The reproducibility of film thickness from run to run as determined by RBS
analysis was better than 5%. The first two samples were grown at $P(\text{GeH}_4)=5\times10^{-4}\text{Torr}$ (as-deposited F-Fast - 6eq. ML/min) and $P(\text{GeH}_4)=2.5\times10^{-4}\text{Torr}$ (as-deposited S-Slow - 3eq. ML/min) in a 10 Torr ambient composed mainly of H$_2$ and immediately cooled to room temperature. The other two samples were grown at the same rate as sample S (reference sample); however, after deposition of the Ge film they were subsequently annealed in-situ for 10 minutes at the growth temperature (600°C) in PH$_3$/H$_2$ with up to $1.4\times10^{-5}$ Torr added PH$_3$ (annealed P) or H$_2$ (annealed H) environments. The samples were characterized initially by Reciprocal-Space Mapping (RSM) using a conventional CuK$\alpha$ X-ray tube and by CVD, at 600°C. Selective chemical etching in 25%NH$_4$OH-31%H$_2$O$_2$ room temperature solution for varying times was used to study the composition profiles in more detail. This etchant is known to be more sensitive to the Ge concentration variation than the RSM measurements, and to slowly remove Ge-rich SiGe alloys with exponentially varying Ge selectivity $^{14,12}$, allowing a detailed study of the remaining material. Although the absolute composition obtained from this technique is not known with a high precision as Anomalous X-ray diffraction $^{2,16}$ or Electron-energy-loss spectroscopy (EELS) in a scanning transmission electron microscope (STEM)$^{17,18}$, the relative comparisons between samples are far more sensitive than either method. For this work our conclusions rely primarily on the relative comparisons rather than on the knowledge of the absolute content. Local and statistical analysis were performed for the as-grown and etched samples with Atomic Force Microscopy (AFM) over ensembles of about 400 islands per etching condition.

Table 1 shows a summary of the growth parameters used and the average Ge composition obtained through the RSM experiments. Samples as-deposited F, as-deposited S and annealed P were found to have the same average Ge content within the experimental uncertainties. However, a lower Ge content was found for sample annealed H, accompanied by a broader diffraction peak corresponding to a wider distribution of compositions within the island ensemble (not shown). These results suggest that Si surface diffusion plays an important role in the final composition profile for sample H, whereas for the other samples that mechanism is minimal. This also confirms the low surface diffusivity of Si in a PH$_3$ environment. The last column displays the total integrated amount of material in the islands, which is consistent with the total amount of Ge deposited of 12.0 ± 0.5ML and the 3.5 ML thick wetting layer.

Figure 1 shows the evolution of samples as-deposited F and S, annealed P and H, before and after etching for 30 min and 60 min. The top images correspond to 250nm × 250nm AFM scans, and the bottom graphs correspond to line profiles taken on statistically representative islands selected from height histograms. The different profiles indicate different degrees of alloying. After 4 hours of etching the entire island material was removed for all cases, leaving a visible moat around the region where the islands were previously located (not shown).

Comparing the AFM images and the line scans before etching, we found that the domes of sample annealed H are slightly larger in diameter; their height is the same as for the dome islands of sample as-deposited S. By comparing samples as-deposited S and annealed P, three main observations can be made for sample P: a) the domes are slightly taller, b) the islands total integrated volume is significantly larger and c) there are no pyramids. From these observations we can conclude that in addition to Ge inter-island diffusion, material from the substrate is effectively being incorporated into the islands causing their growth (comparing samples S and P this amounts to roughly 4 ± 3 ML). This has been observed recently also by STEM-EELS experiments in samples annealed at 650°C$^{13}$, producing a non-abrupt yet uniform interface.

All but sample annealed H consisted of a symmetric Ge-rich outer shell as shown previously by Grazing Incidence Anomalous X-Ray Diffraction (GIXD) experiments$^2$. For sample H an irregular composition profile is revealed by the selective etching, as seen in Figure 2. This has been observed previously in CVD grown samples$^{19}$, and more recently for MBE grown material$^{14}$, and both are consistent with a a significant amount of Si surface diffusion and concomitant intermixing. This asymmetric alloying profile is in accord with the GIXD results of wider composition range within the islands. For sample H, Si and Ge surface diffusion (mechanism b) and possibly intra-island diffusion (mechanism c) act together producing the observed morphology.

For all etching times, sample as-deposited F exhibited a larger fraction of removed material, indicating that intermixing during growth (mechanism a) as well as the other mechanisms are reduced by the shorter growth time. A recent report$^{17}$ on MBE grown material portrayed similar results.

After 30 minutes of etching, the domes in sample as-
deposited S have a top that is richer in Ge compared to sample annealed P. After 60 minutes of etching, one finds that more material has been removed from sample P in contrast to sample as-deposited S and annealed H, indicating a Ge rich region in sample P. These facts demonstrate that Ge and Si redistribute inside the island during annealing in PH$_3$, with Ge diffusing down towards the island base, and Si diffusing up towards the island apex. It is interesting to compare the inferred profile to results of Monte Carlo simulations, which allow for intra-island diffusion$^{[20, 21]}$; both experiment and simulation show a SiGe core and a Ge rich shell.

Summarizing the above results, height statistics for samples as grown and after etching for 30 and 60 min are shown in Figure 2. For samples F, S, H and P the island heights are $12.4 \pm 2.1$ nm, $14.3 \pm 2.3$ nm, $14.1 \pm 2.8$ nm, and $16.1 \pm 2.3$ nm, respectively. The error bars correspond to the standard deviation $\sigma$ of the island height distributions. The line scans (bottom of Figure 1), together with the height statistics, show that for the top 7-8 nm samples annealed P and H are richer in Si when compared to samples as-deposited S and F. This situation is different for the bottom 5 nm, where samples annealed P and as-deposited F exhibit a Ge-rich base. From the evolution of $\sigma$ with etching, one finds that sample annealed H has a consistently broader dome height distribution than the others, indicating a wider composition range within the island ensemble.

The sequence of line scans and height statistics shows that the etching rate is not uniform, indicating that the Ge concentration in the island is not constant in the growth direction. In Figure 3, the total integrated volume is shown as a function of the etching time. The slope of the curve is associated with the average Ge content in the film, with more material being removed faster for Ge rich SiGe regions. Therefore, the Ge content increases from sample H to S, to P and finally F. Comparing samples as-deposited S and F shows that faster Ge deposition results in an increase of the Ge content in the islands, corresponding to less intermixing during growth$^{[17]}$. While sample annealed H is richer in Si compared to the reference sample (as-deposited S), annealing in PH$_3$ enriches the Ge content compared to sample S. This indicates that surface diffusion processes can be selectively controlled depending on the proper choice of the annealing environment.

To understand the intra-island diffusion contribution to the alloy formation, a more detailed analysis was carried out for sample annealed P, where this particular process could be more clearly isolated. The composition profile was studied by sequentially etching and taking line scans along the [110] direction on the same representative island. Figure 4(a) shows line scans for an island before etching and after etching to $h(x, y) < 5$ nm. At this particular height, sample P exhibited a Ge content higher than all samples except sample as-deposited F, indicating Ge diffusion towards the island base (intra-island diffusion). The region after etching shows a smaller diameter and height than prior to etching, indicating a Ge rich shell.

![FIG. 1: AFM images and line scans (bottom) of representative dome islands of all samples as grown, after 30 min and after 60 min of etching with 25%NH$_4$OH:31%H$_2$O$_2$. All images were taken along [110] direction.](image1)

![FIG. 2: Height statistics as a function of etching time for all samples.](image2)
FIG. 3: Total integrated volume (%) as a function of the etching time (minutes) for all samples. The slope of the curve is associated with the average Ge content in the film, with material being removed faster for Ge rich SiGe regions.

It also shows a small dip at the center, corresponding to a Ge rich apex, similar to low temperature MC simulations [21]. In Figure 4(b), a 3D AFM image exhibits a statistically significant rosette pattern, produced by atom redistribution within the island. This observation is associated with strain-assisted intra-island diffusion. Similar morphology has been previously reported, also in a post-growth annealing experiment [3], and was attributed to enhanced surface diffusion, which occurred at the island edges. However, in the present work, alloying takes place inside the island, and the edges remain Ge rich, as can be seen in the line profiles of Figure 4(a); thus no Si could have come from the surface. The driving forces in both experiments are basically the same - minimization of elastic energy and maximization of entropy [21, 22].

The Si-rich regions occur along the [110] directions. Since the facet angle of the bounding \{311\} facets is less steep than the \{153 23\} facets (25° and 32°, respectively) a smaller strain relaxation can take place in the [110] direction. Therefore, Ge moves towards the soft [010] directions and Si moves to the [110] direction, thus producing the rosette structure. At the center of the rosette, one also finds a Ge rich region, presumably reflecting more efficient relaxation at that site. The enhanced intra-island diffusion occurs only very close to the substrate (about 4 nm from the surface of an otherwise pristine island 16 nm high), where the strain is large [22].

In summary, a systematic study focused on the intermixing mechanisms in Ge:Si(001) islands was carried out. Samples were grown and annealed in different environments allowing different diffusion processes to dominate. Selective etching and the RSM experiments permitted a semi-quantitative picture of the Ge content profile inside the dome islands. Increasing the growth rate decreased the degree of Si-Ge intermixing. Intra-island diffusion occurred during annealing in different environments, but surface diffusion could be varied selectively by control-

FIG. 4: Sample annealed P: (a) Line scans on the same island as grown and after two etching steps. (b) 3D AFM image of one representative dome after 60 minutes of etching, showing the rosette final morphology.

ling the ambient gas. When Si surface diffusion was minimized, atomic rearrangement took place within the islands via intra-island diffusion, leading to a four-fold symmetric rosette structure.

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