Secondary ion mass spectrometry characterization of the diffusion properties of 17 elements implanted into silicon

H. Francois-Saint-Cyr
Advanced Materials Processing and Analysis Center, Department of Mechanical, Materials, and Aerospace Engineering, School of Optics, University of Central Florida, Orlando, Florida 32816

E. Anoshkina
Advanced Materials Processing and Analysis Center, Department of Mechanical, Materials, and Aerospace Engineering, University of Central Florida, Orlando, Florida 32816

F. Stevie
Cirent Semiconductor (Agere Systems), 9333 South John Young Parkway, Orlando, Florida 32819

L. Chow
Advanced Materials Processing and Analysis Center, Department of Physics, University of Central Florida, Orlando, Florida 32816

K. Richardson
Advanced Materials Processing and Analysis Center, Department of Physics, Department of Chemistry, University of Central Florida, Orlando, Florida 32816

D. Zhou
Advanced Materials Processing and Analysis Center, Department of Mechanical, Materials, and Aerospace Engineering, University of Central Florida, Orlando, Florida 32816

(Received 26 October 1999; accepted 2 July 2001)

A systematic investigation of the diffusion of Be, B, Na, Mg, Cl, K, Ca, Ti, V, Cr, Mn, Fe, Ni, Zn, Ge, Rb, and Mo in silicon has been carried out. The elements were implanted into silicon wafers as low dose impurities, and then postheat treatments of the ion-implanted samples were conducted at different temperatures for a specific time. Following the anneals, the depth profiles were obtained by secondary ion mass spectrometry analyses. A wide range of diffusion behavior has been observed for these elements. Based on differences in the depth profiles the diffusion mechanism was identified where possible. © 2001 American Vacuum Society. [DOI: 10.1116/1.1396638]

I. INTRODUCTION

The modern semiconductor industry is based on integrated circuits formed from silicon doped with n- and p-type dopants. Diffusion and ion implantation are the two main techniques for introducing different dopants into Si wafers. Since the performance of a device depends critically on dopant distribution, diffusion plays a major role in transistor fabrication. Semiconductor failures are often caused by the diffusion of metallic elements (or contaminants) introduced during device processing. Therefore, a better understanding of diffusion phenomena of metallic elements in silicon is valuable for quality control and failure analysis.

Intrinsic diffusion of metallic impurities in silicon has been under study since the early 1960s.1–4 Most of the data were obtained at that time by annealing a deposited layer on the silicon surface (see Table I). It was noted that, for most elements, the Arrhenius expression is obeyed, indicating thermally activated diffusion. The data compiled show that diffusion is either interstitial or substitutional and there is a wide range of diffusivity for different impurities (over ten orders of magnitude at a given temperature.)

Since the early 1970s, ion implantation5 has been gradually developed into a standard tool to introduce dopants into silicon. Ion implantation generates additional point defects inside Si that further complicate the impurity diffusion phenomena. It is recognized that diffusion of the implanted species during annealing can be affected by the defects generated during the implantation process, in the extreme case causing the formation of an amorphous layer, and by the charge state of the implants. The diffusion behavior of common dopants implanted into Si such as B,6 As,7 Sb,8 and P9 have been well characterized. However, diffusion data on contaminant elements implanted into Si are not generally available. Studies have been made of implanted Fe in InGaAsP10 and Fe and Co in InP.11 In this article, we examine the diffusion behavior of 17 elements, including Be, B, Na, Mg, Cl, K, Ca, Ti, V, Cr, Mn, Fe, Ni, Zn, Ge, Rb, and Mo, implanted into Si.

II. EXPERIMENTAL CONDITIONS

Isotopes of the 17 elements were introduced into (100) single-crystalline silicon by ion implantation at room temperature.51V was implanted into both silicon and a 0.5 μm polysilicon layer on silicon substrate. The implantation energies and doses ranged from 50 to 180 keV and 7.5E13 to 1.1E15 atoms/cm², respectively. The implanted doses for elements with atomic number greater than Si were determined
Focused primary beam was rastered over a 200 μm area, with detection of ions from an area of 60 μm diameter at the center of the raster. The sputtering rate was measured using implanter current comparison with a CAMECA IMS-6f proving Faraday cup and checked by sputtering rate and primary current comparison with a CAMECA IMS-3f using 150 nA primary beam at a source potential of 10 kV, an impact energy of 10 keV, and an impact angle of 40° from normal. The depth scale was established for each profile by measuring the crater depth with a stylus profilometer (Sloan Dektak). The diffusion behavior of the 17 elements in silicon may be categorized into five groups. Since the diffusion behavior of implanted boron in silicon has been extensively studied, it has been used as a control element (category 1). The concentration scales were determined from the known implant doses of the as-implanted samples and the measured erosion rate.

### III. RESULTS AND DISCUSSION

The diffusion behavior of the 17 elements in silicon may be categorized into five groups. Since the diffusion behavior of implanted boron in silicon has been extensively studied, it has been used as a control element (category 1). The concentration scales were determined from the known implant doses of the as-implanted samples and the measured erosion rate.

#### Table I. Diffusion properties of elements in silicon.

<table>
<thead>
<tr>
<th>Element</th>
<th>Ionic radius (nm)</th>
<th>Crystal structure</th>
<th>Diffusion mechanism</th>
<th>Diffusivity at 1000 °C (cm²/s)</th>
<th>Solubility at 1000 °C (atoms/cm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Be</td>
<td>0.034</td>
<td>hcp</td>
<td>I</td>
<td>1.0×10⁻¹⁴a</td>
<td>10²b</td>
</tr>
<tr>
<td>B</td>
<td>0.02</td>
<td>Rhombohedral</td>
<td>S</td>
<td>1.6×10⁻⁶a</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Na</td>
<td>0.098</td>
<td>bcc</td>
<td>I</td>
<td>4×10⁻⁷a</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Mg</td>
<td>0.181</td>
<td>fcc</td>
<td>I</td>
<td>6×10⁻¹⁰a</td>
<td>10⁴b</td>
</tr>
<tr>
<td>K</td>
<td>0.133</td>
<td>bcc</td>
<td>I</td>
<td>2.1×10⁻⁶b</td>
<td>3×10⁴a</td>
</tr>
<tr>
<td>Ca</td>
<td>0.106</td>
<td>fcc</td>
<td>I</td>
<td>2×10⁻⁶b</td>
<td>3×10⁴b</td>
</tr>
<tr>
<td>Ti</td>
<td>0.076</td>
<td>hcp</td>
<td>I</td>
<td>2.2×10⁻⁶b</td>
<td>3×10⁴b</td>
</tr>
<tr>
<td>V</td>
<td>0.065</td>
<td>bcc</td>
<td>I</td>
<td>2.8×10⁻⁶b</td>
<td>5×10⁴b</td>
</tr>
<tr>
<td>Mn</td>
<td>0.091</td>
<td>Cubic</td>
<td>I</td>
<td>2.9×10⁻⁶b</td>
<td>6×10⁴b</td>
</tr>
<tr>
<td>Fe</td>
<td>0.087</td>
<td>bcc</td>
<td>I</td>
<td>7.4×10⁻⁶b</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Ni</td>
<td>0.078</td>
<td>fcc</td>
<td>I/S</td>
<td>10⁻⁷</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Zn</td>
<td>0.083</td>
<td>hcp</td>
<td>I/S</td>
<td>1.0×10⁻⁷</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Ge</td>
<td>0.044</td>
<td>Diamond</td>
<td>S</td>
<td>1.0×10⁻⁷</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Rb</td>
<td>0.149</td>
<td>bcc</td>
<td>I</td>
<td>1.0×10⁻⁷</td>
<td>10⁴b</td>
</tr>
<tr>
<td>Mo</td>
<td>0.068</td>
<td>bcc</td>
<td>I</td>
<td>1.0×10⁻⁷</td>
<td>10⁴b</td>
</tr>
</tbody>
</table>

I—Interstitial.  
S—Substitutional.

*e* 1100 °C.

#### Fig. 1. SIMS depth profiles of ¹¹B (50 keV, 1E15 atoms/cm²) • as implanted, ○ annealed 300 °C, ▲ 500 °C, △ 700 °C, × 900 °C, □ 1100 °C.

#### Fig. 2. SIMS depth profiles of (a) ⁳⁹K (100 keV, 1E14 atoms/cm²) and (b) ⁸⁵Rb (180 keV, 7.5E13 atoms/cm²) • as implanted, ○ annealed 300 °C, ▲ 500 °C, △ 700 °C, × 900 °C, □ 1100 °C for K and 1000 °C for Rb.
500 °C. Diffusion of boron in silicon is first observed for the 700 °C annealed sample. For the samples annealed at 900 and 1000 °C, diffusion of boron occurs both to the surface and deep into the silicon matrix. The second category of the implanted elements shows almost no diffusion after the anneals. Among the elements investigated here, K and Rb have this behavior (see Fig. 2), and Ge, V, and Cl (Fig. 3) show limited diffusion deeper into the silicon matrix (Ge and Cl) or toward the peak of the implant (V). The third category includes Be, Na, and Mg shown in Fig. 4, and Ti and Mn shown in Fig. 5. These elements tend to diffuse to the surface of single-crystalline Si during the anneals greater than 700 °C. Note that significant Ti diffusion to the surface, attributed to interstitial motion, occurs only at 1100 °C.

The fourth category includes the elements Fe, Ca, Cr, Zn, and Mo, and the results are influenced by the defects introduced by ion implantation. It is well known that the ion implantation process will generate a high density of defects in Si and can amorphize the Si crystal at sufficiently high dose. The interactions of the defects and the diffused atoms and the recrystallization of the amorphous Si layer will have a strong effect on the diffusion behavior of the implanted elements. During the annealing process, precipitation of the supersaturated solid solutions and intermediate compounds may occur at the structural defects. The SIMS depth profiles of Fe in the as-implanted and annealed samples, shown in Fig. 6(a), indicate that the implanted Fe atoms in silicon tend to move to the silicon surface during the annealing process. The diffusion of Fe starts for the sample annealed at 500 °C and concentration peaks have been formed at the surface and at 0.13 μm, but the distribution at depths exceeding 0.13 μm was not affected. As the annealing temperature was increased to 900 °C, the implanted Fe atoms diffuse to the silicon surface and form a concentration peak at about 0.02 μm. At this annealing temperature the deeper concentration peak has disappeared. Diffusion of the implanted Cr toward the surface of silicon starts at 500 °C [see Fig. 6(b)], similar
to Fe. Two concentration peaks, located at 0.06 and 0.18 μm, have been observed for the samples annealed at 500, 700, and 900 °C. Most of the implanted dose remains even after significant diffusion has taken place. For example, at 500 °C, 90% of the Cr implanted dose is present, with 5% in the 0.18 μm peak. Notice that the implanted Cr atoms at depths exceeding 0.18 μm are only slightly affected by the 700 and 900 °C anneals. When the annealing temperature was increased to 1100 °C, all the implanted Cr atoms diffuse to the surface. Figure 6 shows that implanted Mo diffuses to the surface of silicon in a manner similar to Fe and Cr, but with different peak locations. This is in agreement with recent work.14

Figure 7(a) presents the SIMS depth profiles of Zn in the as-implanted and the annealed samples, showing that appreciable diffusion of Zn into silicon occurs at 300 °C. Concentration peaks just below the surface and at 0.28 μm have been formed for the samples annealed at 500 or 700 °C. If the annealing temperature is higher than 900 °C, the implanted Zn atoms diffuse to the surface of the silicon. Precipitates at the depth of maximum implant damage and the original amorphous-crystalline interface may explain the existence of the two peaks on either side of the as-implanted profile. The diffusion of Ca, shown in Fig. 7(b), to the surface and into the silicon occurs when the samples were annealed above 700 °C with concentration peaks at 0.1 and 0.15 μm. Concentration peaks at 0.08, 0.17, and 0.27 μm are found for the samples annealed at 900 or 1100 °C.

When three concentration peaks are detected, the first is at the sample surface. The formation of this peak may be explained by the precipitation of solid solutions due to surface defects formed as a result of ion implantation. The second peak occurs at a depth before the projected range, which is the most probable penetration depth for the implant. For example, the projected range for the Cr ion implant is 0.104 μm and the second peak occurs at 0.06 μm. This concentration peak could be due to the formation of compounds and solid solutions. The third peak is deeper than the projected range and is related to end of range implantation defects and, when the dose is sufficiently high, the interface between amorphous and damaged silicon.

Ni in silicon represents the fifth diffusion category. Diffusion of Ni to both the surface and the silicon substrate occurs at 300 °C (see Fig. 8). Nickel has a uniform distribution to 0.2 μm followed by one concentration peak. At 500 °C, two concentration peaks are observed. Cross sectional transmission electron microscopy characterization confirms the precipitation of nickel and the recrystallization of amorphous silicon underneath the surface during the annealing process at high temperatures.

Fig. 5. SIMS depth profiles of (a) Ti (150 keV, 1E14 atoms/cm²), and (b) Mn (150 keV, 1E14 atoms/cm²) [• as implanted, ○ annealed 300 °C, ▲ 500 °C, △ 700 °C, × 900 °C, □ 1100 °C].

Fig. 6. SIMS depth profiles of (a) Fe (100 keV, 1E15 atoms/cm²), (b) Cr (120 keV, 1.1E15 atoms/cm²), and (c) Mo (180 keV, 8.8E14 atoms/cm²) [• as implanted, ○ annealed 300 °C, ▲ 500 °C, △ 700 °C, × 900 °C, □ 1000 °C for Fe and 1100 °C for Cr and Mo].
The mechanism of impurity diffusion in polycrystalline silicon can be quite different from that in single-crystalline silicon. A separate category of diffusion is represented by V in polycrystalline silicon. Figure 9 shows the distribution of V with increasing temperature becomes more concentrated at the interface of the 0.5-μm-thick polycrystalline silicon film and the silicon substrate. In contrast to crystalline Si, diffusion of V in polycrystalline silicon starts at 500 °C and, instead of moving to the surface, the implanted V atoms tend to diffuse to the interface between the polycrystalline silicon and the silicon substrate.

IV. SUMMARY

The diffusion properties of the elements studied can be separated into several categories. For Cl, K, V, Ge, and Rb, very little diffusion has been identified from the SIMS depth profiles. Fast diffusion to the Si surface occurs for Be, Na, Mg, Ti, and Mn if the annealing temperature is above 500 °C. Fe, Ca, Cr, Zn, and Mo show a multi-peaked structure after anneal as low as 500 °C. Ni indicates that it may react with Si and form a layer of silicide in the near-surface region. Finally, the diffusion behavior of V in polycrystalline Si is faster than in single crystalline Si. Diffusion measurements for other elements in Si and a study of the effect of implantation dose are in progress.

ACKNOWLEDGMENTS

The authors would like to thank the other members of the graduate class who participated in this project: B. Prenitzer, B. Kempshall, T. Kimble, B. Surdik, S. Schwarz, S. Skulkarni, C. Urbanik, E. Houge, A. Hussain, and N. Ravikar. The authors acknowledge the technical and financial support from UCF/Cirent Semiconductor Materials Characterization Facility (MCF) and the Advanced Materials Processing and Analysis Center (AMPAC) at the University of Central Florida. The authors also would like to thank R. Irwin, J. Benton, and H. Gossman for valuable discussions, and Zia Ur Rahman, Jay Bieber, and Kirk Scammon for their assistance.
