

Germanium-induced Modulation of Work Function and Impurity Segregation Effect in Fully-Ni-germanosilicide ($\text{Ni}(\text{Si}_{1-x}\text{Ge}_x)$) Gate

Yoshinori Tsuchiya, Atsuhiko Kinoshita, and Junji Koga

Advanced LSI Technology Laboratory, Toshiba Corp.

8, Shinsugita-cho, Isogo-ku, Yokohama 235-8522, Japan.

TEL: +81-45-770-3521 FAX: +81-45-770-3578 E-mail: yoshinori3.tsuchiya@toshiba.co.jp

1. Introduction

It has been recognized that reduction of sheet resistance of gate electrode and suppression of gate depletion are key factors for achieving high-performance MOSFETs in deep sub-50 nm gate regime. Metal gate technology is the solution for these issues. The most difficult challenge is work function control to maintain low V_{th} of less than 0.4 V. Fully-silicide gate is one of the promising candidates for realizing dual work function by segregating impurities at silicide gate electrode/gate insulator interface [1,2]. However, the maximum modulation range ever reported is only 0.32 eV, obtained by Sb, which is insufficient for CMOS application to achieve a low V_{th} .

In this paper, a new method for modulating flatband voltage (V_{fb}) is reported. The effect of Ge incorporation into fully-Ni silicide gate is investigated. It is demonstrated that $\text{Ni}(\text{Si}_{1-x}\text{Ge}_x)$ changes V_{fb} according to Ge fraction: x . In addition, the Ge incorporation modulates the V_{fb} shift effect of B impurity to the opposite direction, compared to the previous report [2]. As a result, the controllable range can be expanded to 0.47 eV near the midgap.

2. Experimental

$\text{Ni}(\text{Si}_{1-x}\text{Ge}_x)$ gated capacitors were fabricated on lightly-doped p-type (100) Si substrates. Following the growth of 10 nm thermally grown SiO_2 dielectric, 50 nm poly-Si or poly-SiGe ($\text{Ge} \leq 30\%$) layers were deposited. Then poly-Si and poly-SiGe were doped with various amounts of B, As and P, and were activated with a 1015°C, 10s anneal. After patterning gate electrode, the gates were silicided with Ni at 450°C or 550°C for 60s in N_2 atmosphere. The CV characteristics were measured at 100 kHz. These samples were analyzed by XRD, TEM, and SIMS.

3. Results and Discussion

A. Effect of Ge incorporation

As shown in Fig. 1, the CV curve shifts to the positive direction, and the work function changes from 4.73 eV to 4.86 eV with increasing Ni thickness. It is identified from the XRD analysis that NiSi is formed in the case of 30 nm Ni sputtered sample, and Ni_2Si is formed in the case of 50 nm Ni sputtered sample (not shown). Therefore, the V_{fb} difference is caused by the phase change from NiSi to Ni_2Si with an increase in Ni-sputtered thickness [3].

In the 30 nm Ni-sputtered sample, a constant V_{fb} value (-0.46 V) is obtained, independent of the Ge fraction (Fig. 2). On the other hand, the V_{fb} values vary with Ge fraction in the case of 50 nm Ni sputtered sample; V_{fb} shifts to the negative direction (-0.42 V) with 5% Ge incorporated, and is fixed until Ge fraction increases to 25%. Then the V_{fb} value shifts to the opposite (positive) direction with 30% Ge incorporated (Fig. 2). As shown in Fig. 3, only in the 50

nm Ni sputtered sample (Fig. 3. (b)), Ni_2Si is formed near the surface region. Near the SiO_2 interface, on the other hand, Ni(SiGe) is formed in both cases. Therefore, the negative shift in the case of $\text{Ge}=5-25\%$ can be caused by the phase change from Ni_2Si to Ni(SiGe) at the SiO_2 interface. From the TEM-EDX analysis, it is identified that Ge can be solved only in the Ni(SiGe) phase. As shown in Fig. 4, in the 50 nm Ni sputtered case, the Ge fraction in the Ni(SiGe) is twice as high as that of poly-SiGe before forming Ni(SiGe), while they are linear in the 30 nm Ni sputtered sample. This is because drive out of Ge from the surface Ni_2Si region results in the Ge condensation in the Ni(SiGe) grain near the SiO_2 interface (Fig. 5). Final Ge fraction in the Ni(SiGe) is about 60% in the 50 nm Ni sputtered sample with an initial Ge fraction of 30%. The positive V_{fb} shift in that highest Ge condensation (60%) (Fig. 2) equivalently agrees with the direction to the work function of NiGe, which is reported as 5.2 eV [4]. This result suggests that higher Ge fraction is required to control the work function between the NiSi and NiGe values.

B. Ge-induced modulation of impurity segregation effect

In the B-doped Ni_2Si samples, V_{fb} shifts to the positive direction, which is similar to the results in Ref. [2]. Inversely, in the B-doped Ni(SiGe) samples, V_{fb} shifts to the negative direction (Fig. 6). The modulation range increases with increasing Ge fraction. The controllable range of work function reaches 0.47 eV by incorporating Ge (4.94 eV ($\text{Ge}=0\%$) to 4.47 eV ($\text{Ge}=30\%$)) (Fig. 7). This value is larger than that of Sb-induced work function modulation [2]. In the NiSiGe sample, Ge is segregated near the gate electrode/ SiO_2 interface (Fig. 8). However, the Ge incorporation has little influence on the segregation of B at the SiO_2 interface, indicating that the impurity segregation effect on work function is affected by the presence of Ge. Although the reason is not clear at present, our experimental results provide a hint for understanding physical mechanism of work function modulation induced by the impurity segregation.

In both As and P implanted samples, V_{fb} shifts to the negative direction as previously reported, and the Ge incorporation expands the modulation ranges (Fig. 9). An addition of the phase change effect (Ni_2Si to Ni(SiGe)) to the impurity segregation effect results in enhanced V_{fb} control range.

4. Conclusion

Ge-induced V_{fb} modulation in fully-germanosilicide gate has been thoroughly investigated. It has been demonstrated that negative V_{fb} shifts, observed in the case of $\text{Ge}=5-25\%$ samples, can be caused by the phase change from Ni_2Si to Ni(SiGe). Higher Ge fraction in Ni(SiGe) ($\text{Ge} \geq 60\%$) is required in order to modulate V_{fb} toward that

of NiGe(positive shift). In addition, the Ge incorporation can expand a controllable V_{fb} range, and the widest range of segregated impurity modulation, which reaches 0.47 eV near the midgap, has been achieved.

Acknowledgment

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Reference

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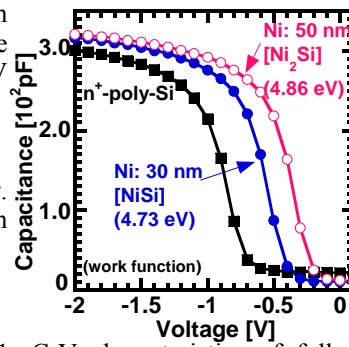


Fig. 1: C-V characteristics of fully-Ni silicide and n^+ -poly Si gates. The values of work function obtained from the V_{fb} -Teff plot are also shown.

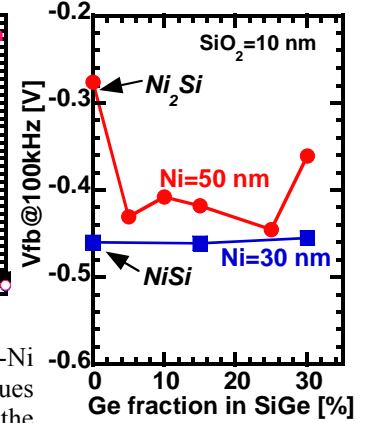


Fig. 2: Dependence of V_{fb} on Ge fraction in SiGe-layer before forming germanosilicide.

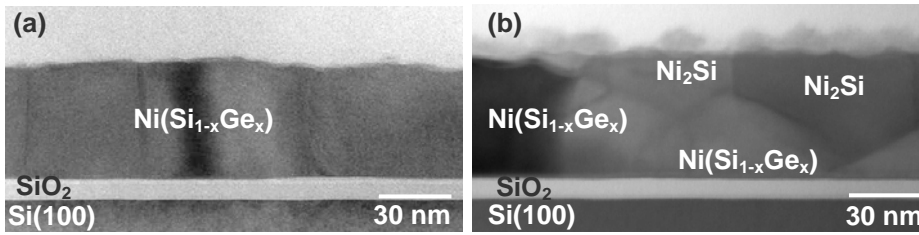


Fig. 3: Cross-sectional TEM image of 15% poly-SiGe layer after forming germanosilicide for (a) 30 nm Ni sputtered and (b) 50 nm Ni sputtered samples, respectively. Ni_2Si is formed near the surface region only in the 50 nm sputtered case.

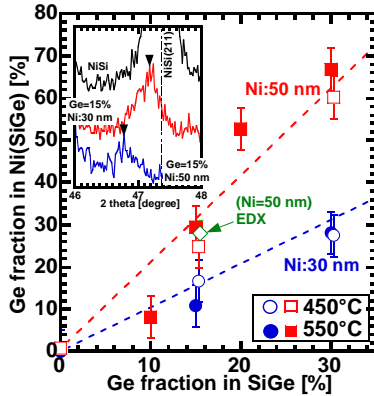


Fig. 4: Relationship between fraction of Ge to Si in Ni(SiGe) and that in poly-SiGe before forming germanosilicide, estimated from XRD Ni(SiGe) diffraction peak shift (insert).

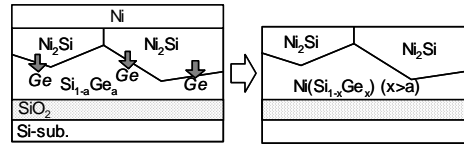


Fig. 5: Schematic of Ge condensation. Drive out of Ge from the surface Ni_2Si region results in the Ge condensation in the Ni(SiGe) grain near the SiO_2 interface.

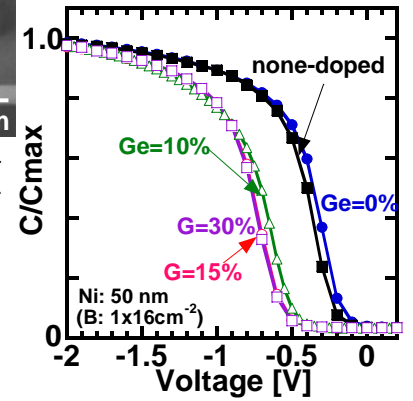


Fig. 6: C-V curves of B-doped samples. Implanted B dose is $1 \times 10^{16} \text{ cm}^{-2}$. In the Ni(SiGe) gates, C-V curves shift to the opposite direction, compared to that in the Ni silicide gates ($Ge=0\%$).

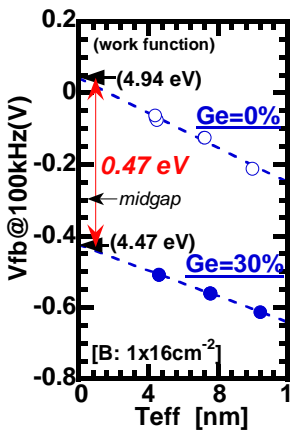


Fig. 7: V_{fb} -Teff plot of B-doped samples.

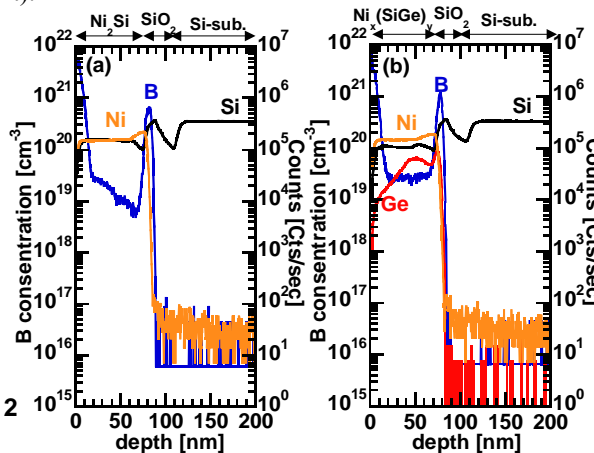


Fig. 8: Backside SIMS depth profile of B-doped (a) Ni_2Si gate and (b) Ni(SiGe) (poly-SiGe with 15% Ge contained) gate. Ge is gathered near the gate electrode/ SiO_2 interface.

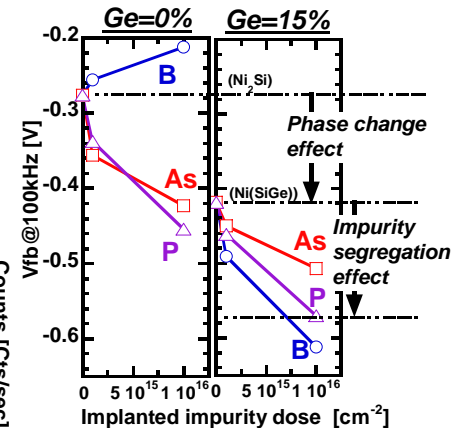


Fig. 9: V_{fb} vs. implanted impurity dose for B, As, and P, in different Ge fractions.