

Formation of C54 TiSi₂ thin films by using high-temperature sputtering and rapid thermal annealing

S. J. LEE, D. Y. KIM

Department of Semiconductor Science, Dongguk University, 3-26 Pildong, Chungku, Seoul 100-715, Korea
E-mail: dykim@dongguk.edu

T. W. KIM

Advanced Semiconductor Research Center, Division of Electrical and Computer Engineering, Hanyang University, 17 Haengdang-dong, Seongdong-gu, Seoul 133-791, Korea

C54 TiSi₂ thin films have become particularly attractive because in comparison with other silicide thin films, they have the unique physical property of lower resistivity with larger thermal stability, as well as a higher possibility of magnetic ordering [1, 2]. C54 TiSi₂ thin films have attracted much attention due to their potential applications in ultra-large-scale integrated devices such as contacts and gate electrodes [3, 4]. Even though several techniques, such as preamorpholization of the Si substrate, growth at high temperature, and insertion of ultra-thin metals [5–8], have been employed to form C54 TiSi₂ thin films, the achievement of high-quality C54 TiSi₂ thin films still requires two-step annealing, which is very complicated. Therefore, the development of a process for the formation of C54 TiSi₂ thin films on p-Si substrates using a one-step method has become very important for device applications. Furthermore, studies on the formation of C54 TiSi₂ thin films using high-temperature sputtering and rapid thermal annealing have not been performed.

This letter reports data on the one-step formation process of C54 TiSi₂ thin films by using a high temperature sputtering and rapid thermal annealing method. X-ray diffraction (XRD) measurements were carried out in order to investigate the phase transitions of the annealed TiSi₂ thin films. X-ray fluorescence (EDX) measurements were performed to characterize the structure and the composition of the TiSi₂ thin films, and sheet resistance measurements were carried out in order to investigate their electrical properties. On the basis of experimental results, a possible formation process is presented for C54 TiSi₂ thin films deposited by using a one-step high-temperature sputtering and rapid thermal annealing method.

The samples used in this study were grown on B-doped p-Si substrates with a (100) orientation. The resistivities of the p-Si substrates at room temperature were approximately between 5 and 10 Ω·cm. The substrates were degreased in trichloroethylene (TCE), rinsed in de-ionized water, etched in a solution of H₂SO₄ and H₂O₂ (4:1) at 40 °C for 5 min, etched in a mixture of HF and H₂O (1:10) at room temperature for 5 min, and rinsed in TCE again. After the chem-

ical process for the Si substrates had been finished, the chemically cleaned wafers were dried at 100 °C in a nitrogen atmosphere for approximately 10 min, and then the substrates were mounted onto a susceptor in the growth chamber. Polycrystalline Ti with a purity of 99.995% was used as the source target material and was precleaned by repeated sublimation. After the chamber had been evacuated to 1×10^{-6} Torr, the deposition was carried out at substrate temperatures of 27, 150, 200, and 250 °C. The rapid thermal annealing process was performed in a nitrogen atmosphere with a tungsten-halogen lamp as the thermal source at temperatures ranging from 450 to 800 °C for various durations.

Fig. 1 shows the XRD curves for titanium-silicide thin films grown on p-Si substrates at 250 °C and subsequently annealed at (a) 450 °C, (b) 500 °C, (c) 550 °C, (d) 600 °C, (e) 650 °C, and (f) 700 °C. While the peak corresponding to the TiSi₂ (210) phase is dominant for the titanium-silicide thin films annealed at 450 °C, the peaks related to the TiSi₂ (210) and the C54 TiSi₂ (040) phase are comparable for the titanium silicide thin films annealed at 500 °C. For the titanium-silicide thin films annealed at 650 °C, the peaks corresponding to the C54 TiSi₂ phase are dominant. Fig. 1 shows that the intensities of the peaks corresponding to C54 TiSi₂ increased significantly with increasing substrate temperature. Therefore, the thickness and the quality of C54 TiSi₂ films are enhanced by increasing both the substrate and the annealing temperatures.

EDX measurements were performed to clarify the chemical components of the thin films after thermal treatment. Fig. 2 shows EDX fluorescence spectra of the TiSi₂ thin films grown on p-Si (100) substrates at (a) 27 and (b) 250 °C and subsequently annealed at 650 °C for 60 s. The results of the EDX measurements show that the grown TiSi₂ thin films consisted of Ti and Si, the stoichiometry of the annealed thin film grown at 27 °C was TiSi_x ($x < 2$), and the ratio between the Ti and the Si compositions of the annealed thin film grown at 250 °C was approximately 1:2, which was consistent, within experimental errors, with stoichiometric TiSi₂. Therefore, EDX measurements revealed that the thin layer grown at 250 °C and annealed at 650 °C for 60 s was TiSi₂.

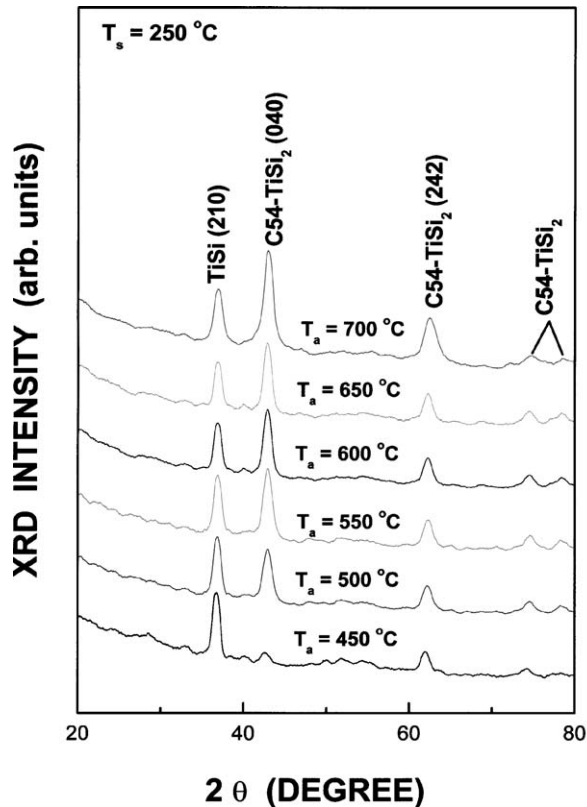


Figure 1 X-ray diffraction curves for titanium-silicide thin films grown on p-Si substrates at 250 °C and subsequently annealed at: (a) 450 °C, (b) 500 °C, (c) 550 °C, (d) 600 °C, (e) 650 °C and (f) 700 °C.

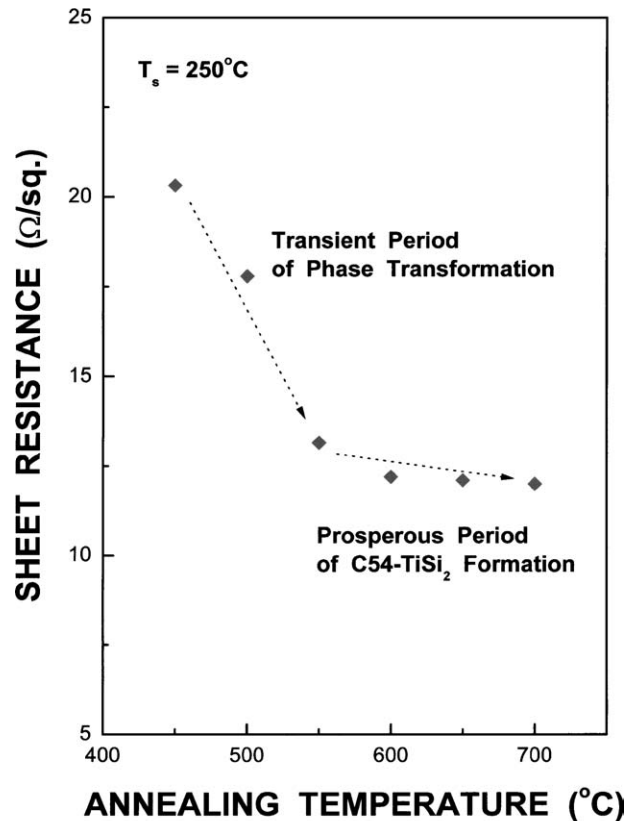


Figure 3 Sheet resistance of titanium silicide thin films grown on p-Si substrates at 250 °C as a function of the annealing temperature.

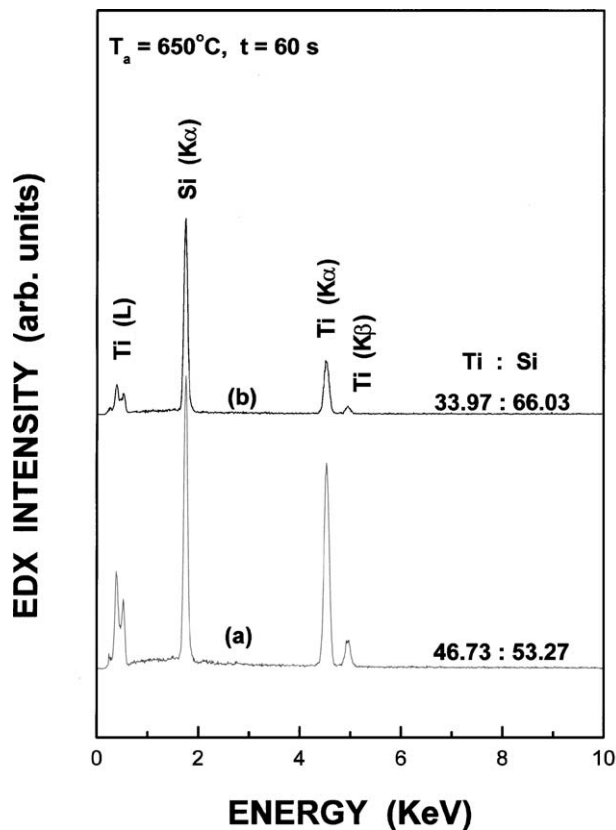


Figure 2 Energy dispersive X-ray fluorescence spectra of titanium-silicide thin films grown on p-Si (100) substrates at: (a) 27 °C and (b) 250 °C, and subsequently annealed at 650 °C for 60 s.

Fig. 3 shows that the sheet resistance of the C54 TiSi₂ thin films grown on p-Si (100) substrates at 250 °C as a function of the annealing temperature. The sheet resistance decreased with increasing annealing temperatures. The dramatic decrease in the sheet resistance at and above 550 °C originated from more C54 TiSi₂ being formed due to a phase transformation from TiSi₂ to C54 TiSi₂. Therefore, increasing of the annealing temperatures reduces the sheet resistance of TiSi₂ thin films.

A possible formation process for C54 TiSi₂ thin films deposited by using high temperature sputtering and rapid thermal annealing can be described on the basis of the XRD results. Even though C54 TiSi₂ thin films are generally formed from C49 TiSi₂ thin films through a two-step process [9, 10], when high-temperature sputtering and rapid thermal annealing is used, the C54 TiSi₂ thin films are formed directly from the TiSi₂ thin films in a one-step process, as shown in Fig. 4.

In summary, C54 TiSi₂ thin films were grown on p-Si (100) substrates by using a high-temperature sputtering method at 270 °C, followed by annealing, and the structural and the electrical properties of the TiSi₂/p-Si (100) heterostructures were investigated using XRD, the EDX, and sheet resistance measurements. The results of the XRD and the EDX measurements showed that the amount of the C54 TiSi₂ formed increased with increasing annealing temperature. The sheet resistance of the C54 TiSi₂ thin films decreased with increasing annealing temperature. A possible one-step formation

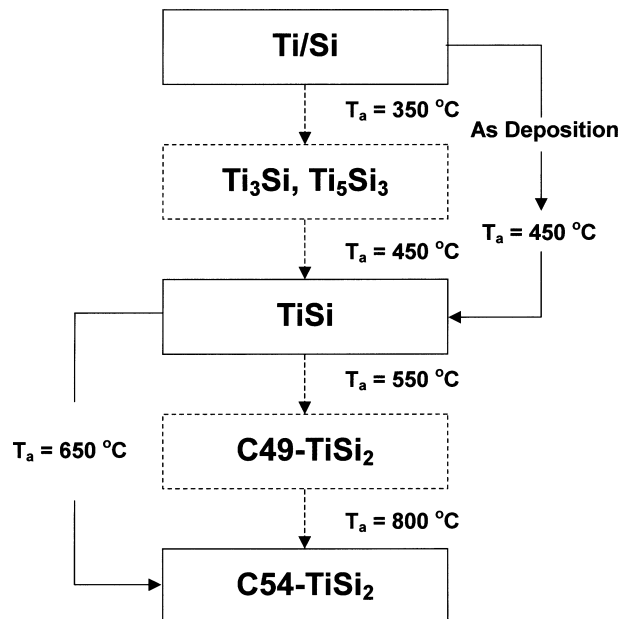


Figure 4 Formation process for C54 TiSi₂ thin films on p-Si substrates. The solid line represents the formation process used in this study for C54 TiSi₂ thin films, and the dashed line indicates the conventional formation process.

process was proposed for C54 TiSi₂ thin films deposited by using high temperature sputtering and rapid thermal annealing.

Acknowledgment

This work was supported by the Dongguk University Research Fund.

References

1. A. V. AMORSOLO, JR. P. D. FUNKENBUSCH and A. M. KADIN, *J. Mater. Res.* **11** (1996) 412.
2. J. A. KITTEL, D. A. PRINSLOW, P. P. APTE and M. F. PAS, *Appl. Phys. Lett.* **67** (1995) 2308.
3. A. MOUROUX, S. L. ZHANG, W. KAPLAN, S. NYGREN, M. OSTLING and C. S. PETERSSON, *ibid.* **69** (1996) 975.
4. S. SINGH, H. SOLAK, N. KRASNOPEROV, F. CERRINA, A. COSSY, J. DIAZ, J. STO and M. SAMANT, *ibid.* **71** (1997) 55.
5. J. A. KITTL and Q. Z. HONG, *Thin Solid Film* **320** (1998) 110.
6. A. LAUWER, A. NAEM, M. DE POTTER and K. MAEX, *ibid.* **320** (1998) 122.
7. A. MOUROUX and S. L. ZHANG, *Appl. Phys. Lett.* **69** (1996) 975.
8. R. T. TUNG, *ibid.* **68** (1996) 1933.
9. S.-L. ZHANG, F. M. D'HEURLE, C. LAVOIE, C. CABRAL, JR. and J. M. E. HARPER, *ibid.* **73** (1998) 312.
10. I. DE WOLF, D. J. HOWARD, A. LAUWERS, K. MAEX and H. E. MAES, *ibid.* **70** (1997) 2262.

Received 7 August

and accepted 24 November 2003