<table>
<thead>
<tr>
<th>Title</th>
<th>Growth and Microstructure of Epitaxial Ti$_3$SiC$_2$ Contact Layers on SiC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Tsukimoto, Susumu; Ito, Kazuhiro; Wang, Zhongchang; Saito, Mitsuhiro; Ikuhara, Yuichi; Murakami, Masanori</td>
</tr>
<tr>
<td>Citation</td>
<td>MATERIALS TRANSACTIONS (2009), 50(5): 1071-1075</td>
</tr>
<tr>
<td>Issue Date</td>
<td>2009-05</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/2433/109943">http://hdl.handle.net/2433/109943</a></td>
</tr>
<tr>
<td>Copyright</td>
<td>© 2009 The Japan Institute of Metals</td>
</tr>
<tr>
<td>Type</td>
<td>Journal Article</td>
</tr>
<tr>
<td>Textversion</td>
<td>Publisher</td>
</tr>
</tbody>
</table>

Kyoto University
Growth and Microstructure of Epitaxial Ti$_3$SiC$_2$ Contact Layers on SiC

Susumu Tsukimoto$^1$, Kazuhiro Ito$^2$, Zhongchang Wang$^1$, Mitsuhiro Saito$^1$, Yuichi Ikuhara$^{1,3}$ and Masanori Murakami$^4$

$^1$World Premier International Research Center, Advanced Institute for Materials Research (WPI-AIMR), Tohoku University, Sendai 980-8577, Japan
$^2$Department of Materials Science and Engineering, Kyoto University, Kyoto 606-8501, Japan
$^3$Institute of Engineering Innovation, The University of Tokyo, Tokyo 113-8656, Japan
$^4$The Ritsumeikan Trust, Kyoto 604-8520, Japan

Growth and microstructure of ternary Ti$_3$SiC$_2$ compound layers on 4H-SiC, which play an important role in formation of TiAl-based ohmic contacts to p-type SiC, were investigated in this study. The Ti$_3$SiC$_2$ layer was fabricated by deposition of Ti/Al contacts (where a slash “/” indicates the deposition sequence) on the 4H-SiC(0001) substrate and subsequent rapid thermal anneal at 1000°C in ultra high vacuum. After annealing, reaction products and microstructure of the Ti$_3$SiC$_2$ layer were investigated by X-ray diffraction analysis and transmission electron microscopy observations in order to understand the growth processes of the Ti$_3$SiC$_2$ layer and determination of the Ti$_3$SiC$_2$/4H-SiC interface structure. The Ti$_3$SiC$_2$ layers with hexagonal plate shape were observed to grow epitaxially on the SiC(0001) surface by anisotropic lateral growth process. The interface was found to have a hetero-epitaxial orientation relationship of $\{0001\}_{\text{TSC}}//\{0001\}_{\text{4H-SiC}}$ and $\langle01T\rangle_{\text{TSC}}//\langle0110\rangle_{\text{4H-SiC}}$, where TSC and 4H-SiC represent Ti$_3$SiC$_2$ and 4H-SiC, respectively, and have well-defined ledge-terrace structures with low density of misfit dislocations due to an extremely low lattice mismatch of 0.4% between Ti$_3$SiC$_2$ and 4H-SiC. [doi:10.2320/matertrans.MC200831]

1. Introduction

Silicon carbide (SiC) is one of the most attractive compound semiconductors for next generation high-power electronic devices operated at high temperature because of its excellent intrinsic properties such as a wide band gap, a high thermal conductivity, a high electric field breakdown strength, and a high saturation electron velocity. However, development of low resistance ohmic contacts to p-type SiC is a key technology issue in the fabrication of highly reliable and high performance devices. Crofton et al. found that binary Ti-Al alloy contacts yielded very low contact resistance to p-type SiC after annealing at temperatures higher than 1000°C. After that, extensive investigations for improvement of the binary Ti-Al contacts and development of other p-type ohmic contact materials have been carried out by various authors.

Microstructures of the TiAl-based ohmic contacts were investigated by transmission electron microscopy (TEM) observations, and these contacts were found to grow Ti$_3$SiC$_2$ on the SiC surface at high temperature of 1000°C in high vacuum. From TEM observations and electrical properties, it is concluded that formation of the Ti$_3$SiC$_2$ layers was required to achieve excellent ohmic properties for p-type SiC, resulting in reduction of height of Schottky barrier formed at the metal/SiC interface and enhancement of carrier transportation across the contact interfaces. However, the role of the Ti$_3$SiC$_2$/SiC interface on mechanism by which the Schottky barrier height reduce has not been well clarified yet. In addition, the atomic structure at/near the Ti$_3$SiC$_2$/SiC interface, which is important to elucidate the carrier transport mechanism, also has not been investigated extensively and understood yet as well as the growth of the Ti$_3$SiC$_2$ compound on SiC has. The primary purpose of the present study is to understand growth process and formation of the Ti$_3$SiC$_2$ compound in the conventional Ti/Al ohmic contacts on SiC annealed at high temperature in order to obtain a clue to reduce the Schottky barrier height. For this purpose, the microstructure at the contact materials/SiC interfaces after isothermal annealing at 1000°C for various storage times are correlated.

2. Experimental Procedures

4H-SiC epitaxial layers (5 μm-thick) doped with aluminum which were grown on undoped 4H-SiC wafers by chemical vapor deposition (manufactured by Cree Research, Inc.) were used as the substrates. Here, the 4H-SiC substrates had 8°-off Si-terminated (0001) surfaces inclined toward a [2110] direction because only 4H-type structure of SiC with polymorph (e.g. 3C, 4H, 6H, 15R etc.) was controllable by lateral growth of the epitaxial layers parallel to (0001)-oriented surface. After chemical cleaning of the substrate surface, a 10 nm-thick sacrificial oxide (SiO$_2$) layer was grown on the SiC substrate by dry-oxidation at 1150°C for 60 min. The substrates were cleaned by dipping in 5% diluted hydrofluoric acid solution and rinsing in deionized water prior to deposition of contact materials. Then, Ti and Al stacking layers with high purities were deposited sequentially on the substrate in a high vacuum chamber where the base pressure was below $5 \times 10^{-6}$ Pa. The thicknesses of the Ti and Al layers investigated in this study are 100 nm and 380 nm, respectively, and these layer thicknesses were chosen to give the average composition of the Ti(20 at%) and Al(80 at%), where the layer thicknesses were measured by a quartz oscillator during deposition. The reasons to choose this average composition was that aluminum rich (more than 75 at%) in TiAl contacts were empirically found to be essential to yield low contact resistance, resulting from formation of the Ti$_3$SiC$_2$ compound layers. After deposit-
ing, the binary Ti/Al contact layers were annealed at 1000°C for various storage times of 2, 4 and 6 min in an ultra-high vacuum chamber where the vacuum pressure was below 1 x 10⁻⁷ Pa.

The surface morphology of the Ti/Al contact layers on 4H-SiC after annealing was observed using a JEOL JSM-6060 scanning electron microscope (SEM). Microstructural analysis and identification of the Ti₂SiC₃ layers at the contact layers/4H-SiC interfaces after annealing was performed using X-ray diffraction (XRD) and cross-sectional TEM. For XRD analysis, Rigaku RINT-2500 with Cu Kα radiation operated at 30 kV and 100 mA was used. In particular, the interfacial structures and an orientation relationship between the contact layers and the 4H-SiC substrates were characterized by cross-sectional high-resolution TEM observations and selected area diffraction pattern (SADP) analysis, respectively, using a JEOL JEM-4000EX electron microscope operated at an accelerating voltage of 400 kV, where the point-to-point resolution of this microscope was approximately 0.17 nm. Thin foil specimens for the TEM observations were prepared by the standard procedures: cutting, gluing, mechanical grinding, dimple polishing, and argon ion sputter thinning techniques.

3. Results and Discussion

3.1 Formation of ternary Ti₂SiC₃ compound on SiC

XRD analysis was performed to characterize the microstructures of the Ti/Al contacts deposited on the SiC substrates after annealing in order to understand growth and formation of the Ti₂SiC₃ compounds. Figures 1 show XRD profiles obtained from the Ti(100 nm)/Al(380 nm) contact layers before and after annealing at 1000°C. Before annealing, the metallic Ti and Al are detected as shown in Fig. 1(a). It is noted that the intensity of the peak diffracted from Ti is very weak because the grain sizes of the as-deposited Ti layer are much smaller than those of the Al layer. After annealing for 2 min, binary Al₃Ti, Al₄C₃, and ternary Ti₂SiC₃ compounds, which were formed by interfacial chemical reaction, are detected by XRD analysis in addition to unreacted Al (Fig. 1(b)). Note that an amount of the Al₄C₃ compounds is quite small because the intensity of the peak diffracted from Al₄C₃ is very weak. After annealing for 4 min (Fig. 1(c)), it is observed that the Ti₂SiC₃ peak intensities increase and the Al₃Ti peak intensities decrease, although the intensities of the peaks diffracted from other compounds do not change significantly. The peaks diffracted from Al₃Ti disappear after annealing for 6 min as shown in Fig. 1(d). From this XRD analysis, only the Ti₂SiC₃ peaks diffracted from (000l) diffraction planes of a hexagonal structure are observed. This result indicates that the Ti₂SiC₃ layer, which is a dominant phase formed by reacting of the Ti-80%Al contact and SiC at 1000°C, has a strong (0001)-oriented texture or hetero-epitaxy on the SiC substrate.

Figure 2 shows an isothermal section of the pseudoternary phase diagram of Al-Ti-SiC system at 1000°C constructed by Viala et al. ²⁹ based on their experimental measurements of Al-Ti-Si-C quaternary system, where a grey region indicates the existence of binary AlTi-based liquid phases, and thick and fine lines indicates sub-solidus lines and tie-lines, respectively. It is noted that the liquidus line would lie at near the composition region of pure Al as denoted by a symbol L in Fig. 2. Based on this phase diagram, the liquid phase is predicted to form when the concentration of Al is more than 75 at% in the Ti/Al contacts at 1000°C. In general, both the solutes and solvents have high diffusivity in liquid, thus the liquid phase has high reactivity to solid (substrate) and the reaction products are easily formed at elevated temperatures. Formation of the liquid is believed to play an important role in formation of the TiAl-based contact by rapid thermal anneal process. Here, the effective average composition of the present reaction system, which react Ti-80%Al and SiC, can assume to be at the point a within region (i) in the diagram. In this region, four phases of SiC, Al₄C₃, Ti₂SiC₃ and liquid with a composition of Al-12%-Si (which is a eutectic composition of Al-Si binary alloy) is predicted to

![Fig. 1 XRD profiles obtained from the Ti(100nm)/Al(380nm) contact layers (a) before and after annealing at 1000°C with various storage times of (b) 2 min, (c) 4 min, and (d) 6 min.](image)

![Fig. 2 Isothermal section of a pseudoternary phase diagram of Al-Ti-SiC system at 1000°C proposed by Viala et al.²⁹](image)
be invariant and coexist in equilibrium. This prediction is consistent with the XRD results obtained from samples annealed at 1000°C for short storage times of 2 and 4 min as shown in Fig. 1(b) and (c), although a small amount of Al<sub>3</sub>Ti and Al did not react completely with SiC and remained at this stage. After annealing for 6 min (as indicated in Fig. 1(d)), disappearance of these unreacted Al<sub>3</sub>Ti and Al is occurred due to additional reaction to form the carbides (Ti<sub>3</sub>SiC<sub>2</sub> and a small amount of Al<sub>4</sub>C<sub>3</sub>) and evaporation of the Al-Si liquid phase with a high vapor pressure during annealing in ultra high vacuum. Thus, the average composition of the reaction system seems to shift toward Al-poor one (region (ii)) as indicated by an arrow and reach to the point b after annealing. In the region (ii), three phases of SiC, Ti<sub>3</sub>SiC<sub>2</sub> and Al-Si liquid with the Si concentrations varied from 12% (at the point c) to 19% (at the point d) are coexisted and their volume will be constituted by an array of tie-lines. However, it is not straightforward to control the compounds formed by rapid thermal annealing process because of occurrence of non-equilibrium phenomena such as evaporation of the liquid phases in this reaction system. This would be a reason why the TiAl-based contact layers have been fabricated empirically without guidelines for designing the materials.

3.2 Growth and microstructure of Ti<sub>3</sub>SiC<sub>2</sub> layers on SiC

In order to characterize the surface morphology of the Ti/Al contact layer after annealing, SEM observation was employed. Figure 3 shows a plan-view SEM image from the Ti/Al contact layer after annealing at 1000°C for 6 min. The surface is observed to have uniformly scale-shaped contrast with hexagonal facet although the surface roughness after annealing was aggravated in comparison with that before annealing (not shown here). Using a stylus surface profiler, the maximum typical roughness was measured to be about 1 µm. The surface facet planes are found to form parallel to (2110) directions on the SiC(0001) substrate surface. The hillocks on the surface due to formation of residual Al-based liquid droplet, forming after annealing at temperatures lower than 800°C, is not observed. This surface morphology results from evaporation of the liquid phases with low melting points and high vapor pressures during annealing at high temperatures in ultra high vacuum as described in the previous section.

Figure 4(a) shows a cross-sectional bright-field TEM micrograph of a typical region in the Ti/Al contact formed on 4H-SiC after annealing at 1000°C for 6 min. The incident electron beam is along the [0110] direction of 4H-SiC, which is parallel to the axis of inclination of the 8°-off SiC (0001) surface. The ternary Ti<sub>3</sub>SiC<sub>2</sub> compound layer forms plate-shaped layer with faceted surfaces. A selected area diffraction pattern at the contact/SiC interface is shown in Fig. 4(b), where the arrays of the diffraction spots from the Ti<sub>3</sub>SiC<sub>2</sub> layer (dotted lines) and 4H-SiC (solid lines) are shown. The Ti<sub>3</sub>SiC<sub>2</sub> layer was found to have an epitaxial orientation relationship with the 4H-SiC substrate:

$$(0001)_{\text{TSC}}/[(0001)_{\text{S}}$$ and [0110]_{TSC}/[0110]_{S}$$

where TSC and S represent Ti<sub>3</sub>SiC<sub>2</sub> and 4H-SiC, respectively. The result of the present TEM investigations is in good agreement with that of the XRD analysis. Both the 4H-SiC (P6<sub>3</sub>/mc; $a = 0.307$ nm, $c = 1.005$ nm) and Ti<sub>3</sub>SiC<sub>2</sub> (P6<sub>3</sub>/mmc; $a = 0.306$ nm, $c = 1.763$ nm) have hexagonal crystal unit cells. The lattice mismatch between the basal planes of Ti<sub>3</sub>SiC<sub>2</sub> and 4H-SiC across the interface is extremely small and approximately 0.4%. The thickness of the Ti<sub>3</sub>SiC<sub>2</sub> layer is not uniform and ranges to 300 nm. The total area of the SiC surface is covered by the Ti<sub>3</sub>SiC<sub>2</sub> layers, meaning that no other compound or reaction layers make direct contact with the SiC surface. In addition, the Ti<sub>3</sub>SiC<sub>2</sub>/SiC interface is observed to have a sawtooth-shaped facet structure as well as the Ti<sub>3</sub>SiC<sub>2</sub> surface is. The (0001)-oriented surface of the Ti<sub>3</sub>SiC<sub>2</sub> layer is also inclined by 8 degrees toward the substrate surface as shown in Fig. 4(a). In the region (ii), Ti<sub>3</sub>SiC<sub>2</sub> layers on SiC layers have a strong influence on the surface orientation of the SiC substrate. Consequently, the scale-shaped surface and interface morphologies is found to originate from growth process and formation of hetero-epitaxial Ti<sub>3</sub>SiC<sub>2</sub> layers on the 8°-inclined SiC(0001) substrate surface based on SEM and TEM observations.

Figure 5 shows a cross-sectional high resolution TEM (HRTEM) image of the interface between the Ti<sub>3</sub>SiC<sub>2</sub> layer and the 4H-SiC substrate, which is taken along the SiC[2110] zone axis. It is observed that lattice fringes run parallel to the interface in both the Ti<sub>3</sub>SiC<sub>2</sub> layer and the SiC substrate. As for the interface morphology, no contamination or second phase layers were observed, and the Ti<sub>3</sub>SiC<sub>2</sub> layer makes direct contact with the SiC substrate. The interface has (0001)-oriented terraces and ledges as marked by letters T<sub>i</sub> ($n = 1, 2, 3, 4$) and L<sub>j</sub> ($j = 1, 2, 3$) in Fig. 5, respectively. The morphology of the terraces is observed to be atomically flat and abrupt. On the other hand, the ledge heights are found to be well-defined and $n \times (a$ half unit cell height of 4H-SiC: 0.5 nm) where $n$ represents the integer, e.g., $n = 11$ for L<sub>1</sub>, $n = 2$ for L<sub>2</sub>, and $n = 1$ for L<sub>3</sub>. This unique interface morphology is caused by chemical reaction of Ti/Al and SiC,
and anisotropic lateral growth of the epitaxial Ti$_3$SiC$_2$ layers along the directions parallel to SiC(0001) as indicated by arrows in Fig. 5. A very small number of misfit dislocations were observed at the interface due to the small lattice mismatch between the Ti$_3$SiC$_2$ and the 4H-SiC substrate as described above. Figures 6 show a cross-sectional HRTEM micrograph magnified the terrace region (T$_4$ in Fig. 5) and a schematic projection of atomic structure at the Ti$_3$SiC$_2$/4H-SiC interface where the unit cell of Ti$_3$SiC$_2$ and 4H-SiC are outlined by rectangle lines. It is noted that the stacking Si-C bilayer marked by termination A of the 4H-SiC substrate surface is observed to make direct contact to Ti$_3$SiC$_2$ at the interface. This is in good agreement with the result reported by Kimoto et al. that the termination surfaces of the A and C bilayers with crystallographically equivalent relation are energetically much stable than that of the B bilayer in 4H-SiC.$^{22}$ From the HRTEM result, the high stability of the termination surfaces A and C in SiC may result in formation of well-defined ledge structure at the Ti$_3$SiC$_2$/4H-SiC interfaces as shown in Fig. 5. In addition, it is apparent that Si monolayer in Ti$_3$SiC$_2$ with layer structure makes direct bond with Si-terminated layer of 4H-SiC(0001) at the
The Si-Si bonding seen intuitively from the present HRTEM image has been suggested by first-principles calculations to be of covalent character. In addition, calculation results (not shown here) also show that the Si-Si interface has smaller adhesion energy compared to other interfaces. The strongest interface predicted by the calculation results (not shown here) also show that the Si-Si interface has smaller adhesion energy compared to other interfaces.

In order to understand chemical reaction processes and growth of ternary Ti$_3$SiC$_2$ compounds fabricated by annealing Ti(100 nm)/Al(300 nm) [total composition: Ti-80%Al] contacts deposited on the 4H-SiC(0001) substrate at high temperature of 1000°C, the microstructures were investigated by XRD analysis and TEM/HRTEM observations. From the XRD analysis, the Ti$_3$SiC$_2$ compound was found to be a dominant phase formed by reacting of the Ti-80%Al contact and SiC at 1000°C, which was in good agreement with prediction of chemical reaction process and products formed during annealing based on thermodynamics (equilibrium phase diagram). The cross-sectional TEM observations revealed that the Ti$_3$SiC$_2$ layers formed by anisotropic lateral growth made direct contact to the SiC substrate surface, and the Ti$_3$SiC$_2$ layers had a hetero-epitaxial relationship with the SiC substrates as follows:

$\{0001\}\overline{[01\overline{1}0]}_{\text{TSC}}//\{0001\}[0\overline{1}0]_{\text{S}}$ (TSC: Ti$_3$SiC$_2$, S: SiC).

Atomically flat terraces and ledges with well-defined heights multiplied about 0.5 nm, which is a half unit cell height of 4H-SiC(0001), were observed at the interfaces between Ti$_3$SiC$_2$ and SiC. Although it seems that the Si monolayer in Ti$_3$SiC$_2$ with layer structure is bonded to the Si-terminated layer of SiC(0001) at the interface, further investigations combined theoretical calculation and atomic-scale characterization are strongly required for determination of the atomic and electronic structures at the interface, which play an important role in formation (carrier transport) of ohmic contacts to SiC.

Acknowledgements

The present work was partially supported by a grant-in-aid for Scientific Research on Priority Area, “Atomic Scale Modification (474)”, from the Ministry of Education, Culture, Sports, Science, and Technology of Japan (No. 19053001).

REFERENCES

20) JCPDS-ICDD PDF No. 22-1317.