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Stable and reliable ohmic contact on p-type 4H-SiC up to 1500 h of aging at 600°C

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Abstract

The stability and reliability at high temperature of Ti_3SiC_2 based ohmic contacts on p-type 4H-SiC (0001) 4°-off substrates were studied. The contact was grown from $Ti_{100-x}Al_x$ alloys annealed at high temperature (from 900°C to 1200°C). The Specific Contact Resistance (SCR) at room temperature and at high temperature (up to 600°C) was in the 10^{-4} - 10^{-5} Ω .cm² range. A Schottky barrier height of 0.71 to 0.85 eV was calculated for the set of samples. After aging period at 600°C for 1500h, the SCR was very stable for Al contents x < 80 at%. This was correlated with chemical and physical stability of these contacts, where the residual stress located on 4H-SiC/ Ti_3SiC_2 interface decreased after aging, for which the Ti_3SiC_2 phase was preserved. Whereas, in the case of x = 80 at%, the Ti_3SiC_2 phase disappeared and the contacts were not ohmic anymore after long time aging. The obtained results showed that $Ti_3SiC_2/4H$ -SiC system is thermodynamically stable at high temperatures and can therefore be a good candidate, with high potential, for high power and high temperature electronic applications.

Keywords:

Silicon carbide, MAX phase, High temperature, Specific contact resistance.

1 Introduction

The semiconductor 4H-SiC is nowadays reaching maturity in both material growth and device performance aspects. Although, current 4H-SiC electronic devices are applied to work at higher temperature than Si- or GaAs-based counterparts, which are limited to temperatures < ~300°C. But several niche applications require device operation at even higher temperatures (> 300°C), such as space (e.g. Venus exploration) or engine monitoring (HT sensors). In fact, the semiconductor 4H-SiC received a high attention by the modern techniques of material growth in high power electronic components because of its special characteristics and abilities in high temperature applications: namely, a good performance at high temperatures; intense powers and high frequencies [1, 2]. 4H-SiC combines exceptional electronic and physical properties [3], such as, a wide bandgap (about 3 times that of Si) [4], high electron saturation velocity and a low intrinsic carrier concentration at high temperatures, comparing to Si, which improves its ability of voltage blocking [5]. Furthermore, 4H-SiC is characterized by its hardness, its low density and its high thermal conductivity [6, 7]. All these properties and much more give 4H-SiC exceptional qualities in high power and high temperature applications.

Among the various reasons for device failure at high temperature is the reliability of the electrical contacts with 4H-SiC [8-11]. Degradation of the contacts usually comes from enhanced chemical reactivity at high temperature. Thus, changes occur in the chemistry of the phases in direct contact with SiC. Ideally, the materials forming the high temperature electric contacts should be in thermodynamic equilibrium with SiC on a wide temperature range to avoid any chemical evolution. On the other hand, the number of metals or materials that can form an ohmic contact on p-type 4H-SiC, is limited, since it is difficult to find metals with an appropriate work function capable of forming an ohmic contact on p-type 4H-SiC which has a high Schottky barrier height (~7 eV) [12, 13]. To obtain such contact on p-type 4H-SiC, aluminum-based alloys are generally used [14]. Many different alloys have been investigated [15], such as Al-Ti [12], Ti/Al/W [13], Ni/Ti/Al [16], Ti/Al/Si [17]. Thierry-Jebali et al. have reported the formation of Ni/Ti/Al contacts on highly p-type 4H-SiC with low Specific Contact Resistance (SCR = $2.8 \times 10^{-6} \ \Omega \cdot \text{cm}^2$), but this result was not reproducible [18]. Another study reported the formation of contacts on highly p-type 4H-SiC after annealing at 800°C (relatively low temperature) for 30 min, but such long annealing time can be detrimental for MOSFET fabrication process [19]. On the other hand, a great attention is given to Al-Ti alloy which has shown stable low SCRs on the order of 10^{-3} – 10^{-5} $\Omega \cdot \text{cm}^2$ on p-type SiC [20, 21]. The formation of the $M_{n+1}AX_n$ phase Ti_3SiC_2 , after annealing of Al–Ti alloy at high temperature, is the reason behind the stability of the ohmic contact on p-type 4H-SiC [12, 22, 23].

The ternary group $M_{n+1}AX_n$ phases (n = 1, 2 or 3) holds its name from its chemical composition, where M is an early transition metal, A is an A-group element (mostly IIIA and IVA) and X is either C and/or N [24-28]. This group of materials combines both metallic and ceramic characteristics: namely, good electrical and thermal conductivity (Metallic properties); good resistance to oxidation and corrosion (Ceramic properties) [29, 30]. Among the MAX phases, Ti₃SiC₂ is a promising material for such high temperature contacts since it is a thermally stable compound which does not react with 4H-SiC at temperatures up to 1200°C and has a Coefficient of Thermal Expansion (CTE) close to that of 4H-SiC [31]. Furthermore, it was shown that this material can lead to low contact resistivity on p-type 4H-SiC [22, 32, 33], which is known to be more difficult than for n-type case. The lowest contact resistivity is obtained when forming Ti₃SiC₂ epitaxial to 4H-SiC (0001) by annealing Al-Ti based alloys [22, 32, 33]. To the best of our knowledge, the high temperature (>300°C) stability and reliability of these contacts have never been investigated. Therefore, an electrical analysis was done using the Transfer Length Method (TLM) to calculate the contact resistivity at high temperatures. In addition, the strain present in the 4H-SiC/Ti₃SiC₂ interface, is estimated with Williamson-Hall analysis (W-H). Afterwards, X-ray Reciprocal Space Mapping (RSM) is applied on 4H-

SiC/Ti₃SiC₂ samples, in order to study the structural properties of the thin film before and after aging, with a particular attention to the epitaxial interface. This is the aim of the present work for which the targeted test temperature was set as high as 600°C.

2 Experimental

2.1 Sample synthesis

P-type doped epitaxial layers (\sim 1.3 μ m thick, [Al] \sim [1-4] \times 10¹⁹ cm⁻³) on 4H-SiC (0001) 4° off substrates were used in this study. The samples were cleaned using acetone and ethanol ultrasonic degreasing, for 5 min each. It was followed by acid treatments: 10 min H₂SO₄:H₂O₂ (2:1) then 4 min HF 5% diluted. Finally, the samples were rinsed with deionized water, and blown dry with N₂ gun. Electrical insulation of the p type layers was done by forming mesa structures using reactive-ion etching (RIE) in an Alcatel Nextral NE110 reactor during 7 min at 60 mtorr under SF₆ (25 sccm) and O₂ (7 sccm) and a RF power of 250 W. After RIE, the samples underwent a second chemical cleaning as mentioned above. Then, 200 nm films of Ti_{100-x}Al_x (50 at% \leq x \leq 80 at%) were deposited at room temperature by magnetron sputtering from three different alloy targets of Ti_{100-x}Al_x composition, with an Ar constant pressure of \sim 5×10⁻³ mbar (Figure 1). The film's composition was determined and checked using Energy Dispersive Spectrometry (EDS).

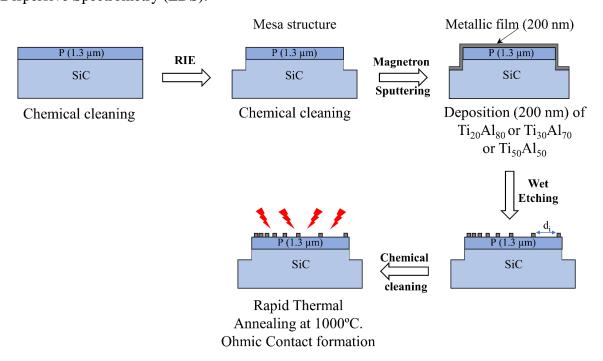


Figure 1: Schematic showing the sample synthesis of Al-Ti-based contact on p-type 4H-SiC, and TLM structure preparation.

2.2 TLM structure preparation

Transfer Length Method (TLM) structures were then fabricated on the mesa, in order to determine the electrical properties of the metal/p-type 4H-SiC contact. This was performed by standard photolithography followed by wet etching using commercial Al-etch at 60° C. The TLM pattern was composed of seven rectangular electrodes ($500 \times 100 \ \mu m^2$) with increasing spacings of 3, 6, 10, 20, 40, 80, and 120 μ m. For ohmic contact formation, rapid thermal

annealing (RTA) of the $Ti_{100-x}Al_x$ alloys was performed under argon at atmospheric pressure. The heating rate was 20°C/s and the annealing plateau was set to 1000°C for 10 min as these conditions were found to give the best results in terms of contact resistivity (SCR = 9.8×10^{-6} $\Omega\cdot\text{cm}^2$ at 600°C , according to ref [32]). After optical microscopy inspection of the annealed TLM patterns, we could confirm that their dimensions and the interpad distances did not change so that no correction was needed during the TLM measurements. Note that the length of the rectangular pads, $500~\mu\text{m}$, is large enough compared to the distance between the pads and to the distance between the pads and the edge of the mesa (5 μm), so that the current crowding effect can be considered negligible.

The structural properties of the contacts were routinely determined by X-ray diffraction (XRD) on a 'Rigaku Smartlab' diffractometer equipped with a Cu rotating anode working at 9 kW, and the wavelength was set to $\lambda = 1.54056$ Å. The electrical characterization was done using a Keithley source measurement unit K2602A, where the samples are mounted directly on a heating chuck of a probe station. They are characterized at different temperature, up to 600°C, and monitored using a K-type thermocouple. The temperature accuracy of the tests was of $\pm 1\%$. Above 300°C, electrical tests are performed under vacuum to prevent oxidation of the TLM structures. A four-probe setup is used (Kelvin configuration) to eliminate cable and tips resistances, which results in a measurement accuracy value better than 1%. I-V characteristics are plotted as a function of the contact spacing of different temperatures.

2.3 Aging conditions

To study the reliability of these contacts, the aging conditions were 600°C under pure argon (not under air, to prevent oxidation of Al at surface) for durations up to 1500 h. These aging tests were carried out in a resistively heated tubular furnace. The evolution of contact nature and resistivity was determined after each aging period intervals (24, 48, 100, 200, 400, 700, 1000 and 1500 h) using the following procedure: 1) the samples were cooled down to room temperature, then 2) they underwent XRD analyses and contact resistivity measurements after which 3) they were put back in the aging furnace and heated again to 600°C for the subsequent aging interval.

3 Results and discussion

3.1 Specific contact resistance

For $Ti_{20}Al_{80}$ —based contact, the evolution of the specific contact resistance (SCR) as a function of aging time is shown in Figure 2 (black points). It was found to decrease from 4.1×10^{-4} to $2.1\times10^{-4}~\Omega.\text{cm}^2$ within the first 48 h of aging. For longer aging time, the I-V characteristics were not linear anymore which means that the contact became non-ohmic. When looking to the XRD patterns recorded before and after 100 h of aging, one note that the ohmicity degradation could be correlated to compositional changes of the metallic phases present in the contact (Figure 3). Indeed, after aging the diffraction peaks corresponding to Al and Ti_3SiC_2 almost disappeared while the intensity of the Al_3Ti peaks increased. As a matter of fact, the ohmicity loss can be correlated to the disappearance of the Ti_3SiC_2 phase.

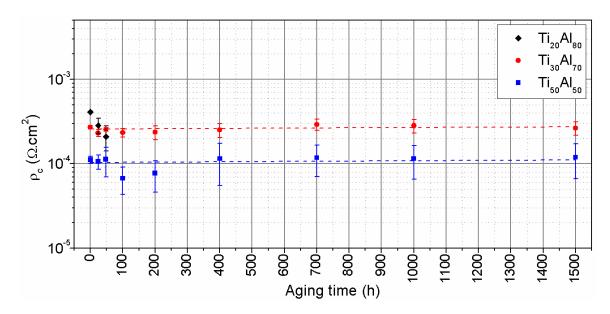


Figure 2: Variation of the SCR as a function of aging time for Ti20Al80, Ti30Al70 and Ti50Al50-based contacts.

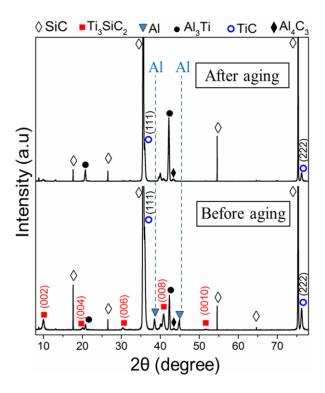


Figure 3: XRD patterns before and after aging of Ti20Al80-based contact.

This is not the case for $Ti_{30}Al_{70}$ and $Ti_{50}Al_{50}$ -based contacts for which the SCR kept constant even after 1500 h of aging at 600° C (Figure 2). When looking to the corresponding XRD patterns of $Ti_{50}Al_{50}$ -based contacts in Figure 4, it is obvious that only the Al peaks disappeared after aging, while the ones of $Ti_{3}SiC_{2}$ were still present, though of lower intensity. Also, less obvious but detectable is the evolution of $Al_{3}Ti$ peaks: they increased while shifting towards low angles. This is better seen in Figure 5, for which the XRD patterns were recorded after only 24 h of aging. In fact, these few changes in the contacts composition seems to happen during

the early aging time (\leq 24 h) while after that no further change is observed. We can therefore conclude that the system reaches a thermodynamic stability in the first aging hours.

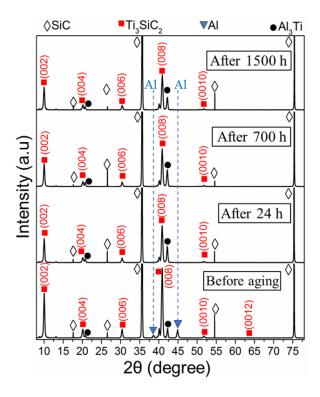


Figure 4: XRD patterns before and after different aging times of Ti₅₀Al₅₀-based contacts.

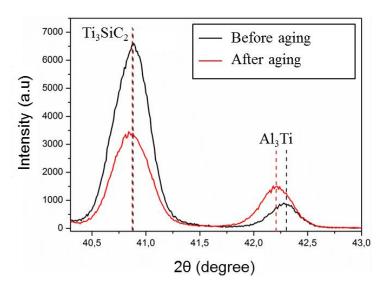


Figure 5: Zoom on the XRD pattern recorded for $Ti_{50}Al_{50}$ -based contact, before and after 24 hours of aging at 600°C. Note after aging the decrease of the Ti_3SiC_2 peak intensity and the increase and shift, to lower angles, of the Al_3Ti peak.

Many investigations focused on the thermal stability of ohmic contacts on p-type 4H-SiC at high operating temperatures up to 600°C. A reported Ni/Ti/Al ohmic contacts on p-type 4H-SiC showed significant changes in the contact resistivities and an electrical degradation after aging at 600°C for 100h [16]. Vivona et al. have reported the thermal stability of Ti/Al/W

contacts to p-type implanted 4H-SiC at 400°C, and the results showed some changes in the contact resistivity after 100h of aging [13]. Furthermore, recent studies presented the thermal stability of Ti/Pt, Ni/Ti/Ni/TaSi₂/Pt and Ni/W/TaSi₂/Pt contacts to p-type implanted 4H-SiC at 500°C for 300h [34, 35]. The SCR of Ti/Pt increased dramatically to 2 Ω ·cm² after 200h of aging at 500°C [34]. In addition, the SCR value of Ni/Ti/Ni/TaSi₂/Pt contact on p-type 4H-SiC increased in the initial 50h of aging at 500°C [34]. Moreover, the reported Ni/W/TaSi₂/Pt ohmic contact on p-type 4H-SiC showed some changes in the SCR on the order of 10^{-3} Ω ·cm² after 300h of aging at 500°C [35]. On the other hand, the results presented in Figure 2, of this study, showed an extreme electrical stability for Ti₃₀Al₇₀ and Ti₅₀Al₅₀-based contacts where the SCR kept constant at 600°C even after 1500h, which is relatively a long time of aging. The obtained results demonstrated the reliability and the thermal and electrical stability of Ti₃₀Al₇₀ and Ti₅₀Al₅₀-based contacts on p-type 4H-SiC, which is promising for high temperature applications.

3.2 Chemical analysis

As a matter of fact, for all the samples of this study, the observed phases evolution after aging is rather similar:

- Pure Al phase disappears
- Ti₃SiC₂ phase amount decreases
- Al₃Ti phase amount increases

Based on these results, we can propose that the following reaction takes place during the aging at 600°C:

$$Ti_3SiC_2 + 9Al \rightarrow 3Al_3Ti + SiC + C$$
 (1)

This reaction is obviously not complete in the case of $Ti_{30}Al_{70}$ and $Ti_{50}Al_{50}$ -based contacts since Ti_3SiC_2 phase is still detected in these samples, even after 1500 h of aging. This is probably due to the exhaustion of elemental Al which stops the reaction. But, such reaction does not explain the low-angle shift of Al_3Ti peak after aging. The literature mentions that Al_3Ti can contain up to 15% at. of Si in its lattice, forming a $(Al-Si)_3Ti$ solid solution [36] and shifting its XRD peaks to low angles. Assuming the formation of such solid solution, we can rewrite equation (1) as follows:

$$Ti_3SiC_2 + 9(1-x)AI \rightarrow 3(Al_{1-x}Si_x)_3Ti + (1-9x)SiC + (1+9x)C$$
 (2)

It is interesting to compare these reactions to the one proposed in [37] for the formation of Ti_3SiC_2 during the $1000^{\circ}C$ rapid thermal annealing:

$$20Al + 9Ti + 2SiC \rightarrow Ti_3SiC_2 + 6Al_3Ti + 2Al + Si$$
 (3)

Obviously, reaction (3) cannot be considered as the reverse of reactions (1) or (2). According to ref [38], annealing an Al-rich Al-Ti alloy on SiC at 700°C leads to the only formation of Al₃Ti phase, almost without Ti₃SiC₂. As a matter of fact, Al₃Ti seems to be the equilibrium phase with SiC at low temperature ($\leq \sim 700$ °C), while this equilibrium shifts toward Ti₃SiC₂ formation at higher temperatures. Assuming this, we can now explain the chemical evolution of the contacts after aging at 600°C: such lower temperature treatment provokes the displacement toward Al₃Ti phase of the equilibrium with SiC, at the expanse of Ti₃SiC₂ phase. Due to some probable loss of Al (by evaporation) during the various annealings, reaction (2) is not complete so that the system does not shift completely to the equilibrium with Al₃Ti and then both Ti₃SiC₂ and Al₃Ti phases coexist. The important consequence of such incomplete reaction is that the contact is in thermodynamic equilibrium with SiC as long as no Al is added to the system. It means that, as long as only chemical reactivity is governing the contacts ohmicity, this contact can virtually remain perfectly stable at 600°C, no matter the aging time.

Finally, note that both reactions (1) and (2) should lead to the formation of SiC and C. Unfortunately, we could not confirm this point since the XRD spectra did not display evidence of these phases. Further analyses are programmed to evidence the possible presence of elemental carbon, such as Raman spectroscopy or cross-section elemental analyses. But in all cases, these extra phases do not seem to be detrimental to the contact stability and thus no visible impact on the overall thermodynamic equilibrium is detected.

3.3 Strain and Stress

In the following, we used the Williamson-Hall (W-H) method to investigate the origin of the X-ray diffraction peaks broadening. It is obvious from equation (5) that the corrected peak broadening β_{hkl} (in radians) varies as $1/\cos\theta$ from crystallite size (L) and as $\tan\theta$ from strain (ϵ). The W-H method consists on separating the size and strain broadening by presenting the peak width as a function of the diffracting angle 2θ , giving the following equation:

$$\beta_{hkl} = \beta_L + \beta_s \tag{4}$$

$$\beta_{hkl} = \frac{k\lambda}{L \cos\theta} + (4\epsilon \tan\theta)$$
 (5)

By rearranging Eq. (5):

$$\beta_{hkl}\cos\theta = \frac{k\lambda}{L} + 4\varepsilon\sin\theta \tag{6}$$

Two graphs were plotted for $Ti_{50}Al_{50}$ -based contacts before and after aging up to 1500h at 600°C, with '4sinθ' defining the x-axis and ' β_{hkl} cosθ' defining the y-axis. The plots are illustrated using the preferred orientation peaks of Ti_3SiC_2 . Afterwards, a linear fit is applied on the plots, and the strain is determined from the slope of the fit. Figure 6 shows the W-H plots for Ti_3SiC_2 -based contacts before and after aging up to 1500h at 600°C. It is remarkable that the strain in the Ti_3SiC_2 thin film before aging is larger than after aging, meaning that aging has an effect on the strain located in the contact. In addition, the stress exerted on the contact can be determined by using the generalized Hooke's law $\sigma = E \cdot \varepsilon$, where the linear proportionality between stress(σ) and strain(ε) is considered, with Young's modulus (E = 326GPa for Ti_3SiC_2 -based contact has reduced from 145MPa to 11MPa which means that the stress in the contact has almost totally relaxed after being subjected to a temperature of 600°C up to 1500h, and no serious deformation or damage have occurred in the physical structure of the contact.

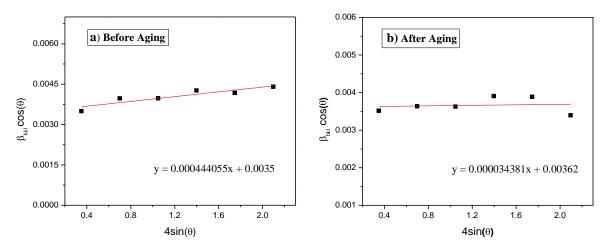


Figure 6: W-H analysis of Ti₅₀Al₅₀-based contacts, a) Before aging, b) After aging up to 1500h at 600°C.

3.4 Reciprocal space mapping

Figure 7 illustrates the RSM patterns around 4H-SiC (1 0 7) and Ti₃SiC₂ (1 0 12), which were used to determine the d-spacing for each phase. Because of the hexagonal structure of 4H-SiC and Ti₃SiC₂, the lattice parameter 'a' was calculated by the following equation [40]:

$$\frac{1}{d^2} = \frac{4}{3} \left(\frac{h^2 + hk + k^2}{a^2} \right) + \frac{l^2}{c^2} \tag{7}$$

The obtained results are presented in Table 1.

Table 1: Lattice parameters.

	Lattice parameter (Å)		
	Theoretical value	Before aging	After aging
4H-SiC (1 0 7)	a = 3.079	a = 3.079	a = 3.079
	c = 10.061	c = 10.061	c = 10.061
Ti ₃ SiC ₂ (1 0 12)	a = 3.068	a = 3.079	a = 3.069
	c = 17.669	c = 17.656	c = 17.667

Table 1 shows that the calculated lattice parameter a = 3.079 Å of 4H-SiC is equal to the theoretical value, which is 3.079 Å and didn't change before and after aging. Besides, it is remarkable that for Ti_3SiC_2 , before aging the parameter 'a' is equal to that of 4H-SiC which means it is larger than the theoretical value a = 3.068 Å. Therefore, due to the difference between the theoretical and the calculated value of the lattice parameter of Ti_3SiC_2 , stress was introduced into the contact. On the other hand, after aging the lattice parameter 'a' of Ti_3SiC_2 reduced (a = 3.069 Å) and reached a closer value to the theoretical one (a = 3.068 Å), leading to a stress relaxation in the Ti_3SiC_2 -based contact.

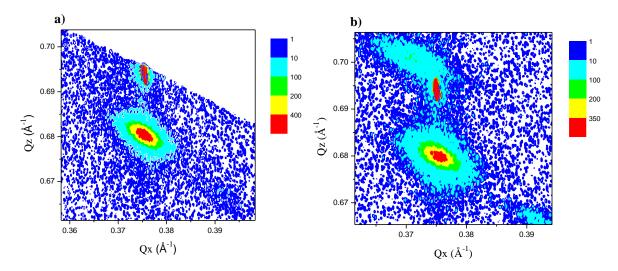


Figure 7: Reciprocal space maps around SiC (1 0 7) and Ti_3SiC_2 (1 0 12), a) Before aging, b) After aging.

In brief, the synthesized layer of Ti_3SiC_2 adopted a lattice parameter equal to that of the 4H-SiC substrate, which caused some strain in the Ti_3SiC_2 layer. However, the aging at 600°C for 1500h permitted the Ti_3SiC_2 layer to reach a lattice parameter closer to the normal one of the Ti_3SiC_2 phase. The obtained results showed that the strain and the stress exerted on Ti_3SiC_2 -based contact reduced after aging, this means there is no serious structural deformation occurring to the Ti_3SiC_2 layer, consequently 4H-SiC/ Ti_3SiC_2 interface is intact. The good mechanical behavior and the physical constancy of the contact could be one of the reasons that improve the reliability of the contact, and leading to a low and stable electrical resistivity even at high temperatures.

Conclusion

In this paper, an electrical study was done using Transfer Length Method (TLM) to calculate the specific contact resistance of Ti₃SiC₂-based contact formed on a 4H-SiC p-type substrate. The TLM was applied on the samples before and after aging for 1500h at 600°C. In addition, strain, stress and lattice parameters located in Ti₃SiC₂ thin films were determined in order to examine the effects of high temperature on the physical structure of the electrical contact and 4H-SiC/Ti₃SiC₂ interface. In conclusion, Al-Ti based metallization on p-type 4H-SiC allows elaborating low resistivity and reliable contacts for operating temperatures as high as 600°C. This can be achieved for an initial Al content in the 50 - 70 at% range which produces the suitable 4H-SiC/Ti₃SiC₂ interface which is not converted back to Al₃Ti upon aging at 600°C for up to 1500h. Besides, mechanical stress, exerted on the electrical contact, decreased after aging, which is leading to a relaxation in the contact and preventing a serious structural deformation. The physical stability helps the contact to keep a steady resistivity, as well as the chemical balance. Such interface is thermodynamically stable and can thus virtually remain perfectly stable and reliable at temperatures up to 600°C.

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