Strain Effect in Highly-Doped n-Type 3C-SiC-on-Glass Substrate for Mechanical Sensors and Mobility Enhancement

Hoang-Phuong Phan,* Tuan-Khoa Nguyen, Toan Dinh, Han-Hao Cheng, Fengwen Mu, Alan Iacopi, Leonie Hold, Dzung Viet Dao, Tadatomo Suga, Debbie G. Senesky, and Nam-Trung Nguyen

This work reports the strain effect on the electrical properties of highly doped n-type single crystalline cubic silicon carbide (3C-SiC) transferred onto a 6-inch glass substrate employing an anodic bonding technique. The experimental data shows high gauge factors of 8.6 in longitudinal direction and 10.5 in transverse direction along the [100] orientation. The piezoresistive effect in the highly doped 3C-SiC film also exhibits an excellent linearity and consistent reproducibility after several bending cycles. The experimental result is in good agreement with the theoretical analysis based on the phenomenon of electron transfer between many valleys in the conduction band of n-type 3C-SiC. Our finding for the large gauge factor in n-type 3C-SiC coupled with the elimination of the current leak to the insulated substrate could pave the way for the development of single crystal SiC-on-glass based MEMS applications.

1. Introduction

The demand for electronics and sensors which can operate in extreme conditions has propelled the research into wide band gap materials such as III-nitride (e.g., GaN, AlN), silicon carbide (SiC) and diamond-like carbon (DLC). Among these materials, SiC has emerged as an excellent semiconductor for Micro Electro Mechanical Systems (MEMS) applications owing to its large energy gap, varying from 2.3 to 3.4 eV, excellent chemical inertness, superior mechanical properties, and outstanding radiation tolerance. Numerous SiC technologies have been transitioned into commercial products, including bipolar transistors, Schottky diodes, field effect transistors, ultraviolet photodetectors, and gas sensors.

Although great progress has been made to commercialize SiC materials and devices, the material cost is still relatively expensive in comparison to the Si counterpart. One promising approach to bring down the cost of SiC materials is the development of epitaxial SiC nanofilm on a Si substrate. This approach can utilize the low cost and worldwide availability of Si substrate, as well as take advantage of Si-orientated MEMS fabrication technologies. To date, high quality and large scale SiC wafers grown on Si have been reported. Applications of SiC-on-Si have also been demonstrated such as mechanical sensors, chemical sensors, bio devices. Nevertheless, when subjected to high temperature, the current leakage from SiC to the Si substrate could adversely affect the performance of the SiC sensing element. Therefore, a number of efforts have been made to transfer SiC films from SiC/Si wafers onto electrically insulating substrates. Most of the works reported in the literature are based on SiO2 to SiO2 direct bonding or Si to poly Si diffusion bonding techniques to form an oxide layer sandwiched between SiC and Si. However, these methods require an ultra-smooth buffer layer such as SiO2, formed by sputtering/oxidation and polishing or SiO2/poly-Si which typically complicates the process. Furthermore, the Si layer in the bonded wafers generally absorbs visible wavelengths, hindering the merit of high-optical-transmittance in bulk SiC materials. Employing the anodic bonding method, we have recently demonstrated large scale single crystal 3C-SiC films transferred onto a glass substrate. The bonded wafers possess
robust adhesion strength, high visible-wavelength transmittance, and excellent electrical insulation to the substrate. In addition, the transferred SiC nanofilm has also been proven to be bio-compatible which enabled the direct culture of bio-cells. Furthermore, the large scale of 6-inch in the diameter also makes the single crystal 3C-SiC/glass wafer a promising platform to develop low-cost mass production devices for a wide range of MEMS applications.

This work elucidates the strain effect on the electrical conductance of highly doped n-type 3C-SiC nanofilms bonded onto glass through experimental characterizations and the first principle calculation. The experimental results show a negative strain gauge factor of $-8.6$, which is in good agreement with theoretical analysis based on the many-valley electron transfer phenomenon. The insight obtained from this study will be the foundation for development of SiC/glass mechanical sensing applications as well as opening up opportunities for further investigation into strain engineering in SiC.

2. Results and Discussion

To conduct this experiment, single-crystalline 3C-SiC thin films were grown on 6-inch Si(100) substrates using a low pressure chemical vapor deposition process (LPCVD). Silane (SiH4) and propene (C3H6) were employed as the precursors in the hot-wall furnace at 1250°C. Additionally, ammonia (NH3) was utilized as the in situ dopant, forming highly-doped n-type 3C-SiC. The thickness of the initial 3C-SiC film was 700 nm, measured using a NANOMETRICS Nanospec/AFT 210. The $\theta - 2\theta$ scan of X-Ray Diffraction (XRD) shows three peaks at 41.4°, 69.1°, and 90°, corresponding to 3C-SiC(200), Si(400), and 3C-SiC(400), respectively. This result indicates that the SiC film was epitaxially deposited on the Si(100) wafer. In addition, the bright spots observed in the selected area electron diffraction (SAED) indicates that the 3C-SiC film was single crystal (See supplementary material). The surface roughness of the as-deposited SiC film was found to be approximately 10 nm using atomic force microscopy. The film was further smoothed by etching back the SiC layer to a thickness of 600 nm using plasma assisted chemical etching. The RMS roughness of the smoothed wafer was found to be less than 0.5 nm as shown in Figure 1(a) which significantly improves the contact surface between SiC and glass during wafer bonding. The detail of the SiC-smoothing process can be found elsewhere.

Prior to the bonding process, piranha cleaning and oxygen plasma was performed to remove contamination from the surface of glass (sourced from UniversityWafer™ with a thickness of 500 μm and surface roughness of less than 1.5 nm) and SiC/Si wafers. The smoothed 6-inch SiC/Si wafers were then subjected to the anodic bonding process, performed using an EVG 520IS hot embosser under a compressive force between 1.5 and 2.5 kN and bias voltage between $-200$ to $-1000$ V at a temperature of 400 °C. In the anodic bonding process, the SiC/Si wafer was connected to anode, while the cathode electrode was applied to the glass wafer. Subsequently, the cathode attracted positive charges from the glass, resulting in a depletion layer at the interface between SiC and glass. As a consequence, the oxide negative charges in the glass moved toward the anode and diffused into the SiC to form chemical bonds.

![Figure 1](https://example.com/figure1.png)

**Figure 1.** Fabrication of SiC/glass piezoresistors. (a) AFM image of the surface of a smoothed SiC film prior to bonding; (b) The concept of anodic bonding process and lithography on the SiC/glass platform; (c) Photograph of a full 6-inch SiC/glass wafer; (d) Photograph of a SiC/glass cantilever used in the bending experiment (Inset: Microscope image of SiC piezoresistors aligned in longitudinal and transverse [100] orientations).
bonding between the SiC and glass substrate. Using Depth profile X-ray photoelectron spectroscopy the bonding layer was found to be less than 15 nm. The bonding strength was characterized using a pulling test, showing a large tensile strength above 20 MPa, which is excellent for mechanical sensing.\(^{[27]}\) Subsequently, the top Si layer with a thickness of 650 \(\mu\)m was completely removed in KOH at 80°C. Since the etching selectivity between Si and glass in KOH is above 500:1, the thickness of the glass layer did not significantly changed after Si removal.\(^{[31]}\) In addition, due to its excellent chemical inertness, the SiC film functioned as an etch-stop layer, preventing KOH from attacking the bonding interface.

A two-mask fabrication process was carried out to fabricate SiC resistors using Inductive Coupled Plasma (ICP) with an etching rate of 100 nm min\(^{-1}\). Our standard etch processes have been developed in the STS\(^{TM}\) ICP plasma etcher that utilizes an electrostatic chuck to provide excellent etch uniformity (<1%) on a 150 mm wafer. By using a very low etch rate process, we can provide a controlled removal of as little as approximately 15 nm at a time. A longer etch time gives a proportionally greater etch depth and the etch depth was measured using Dektak 150, confirming that the 600 nm SiC was completely etched through to the glass substrate.

A nickel film with a thickness of 300 nm was then deposited using sputtering and then annealed at 500°C to form the contact to SiC resistors. Finally, the SiC on glass samples were diced into cantilevers with dimensions of 0.51 \(\times\) 3.5 \(\times\) 20 mm\(^3\) for the subsequent bending experiment, as illustrated in Figure 1(b).

Figure 1 shows a 6-inch 3C-SiC/glass wafer, exhibiting excellent transparency. The optical characterization and the bonding strength of the films can be found elsewhere.\(^{[27]}\) The large scale of the bonded wafers could significantly reduce the material cost as well as allow mass production of SiC/glass based devices by employing a batch fabrication process. Figure 1(d) shows a photograph of SiC/glass cantilever for the bending experiment, in which the U-shaped SiC piezoresistors with dimensions of 8 mm \(\times\) 200 mm \(\times\) 600 mm were patterned near the fixed end of the cantilever to obtain high tensile strain.

The semiconductor type and the carrier concentration of the bonded 3C-SiC films were characterized using a hot probe technique.\(^{[30]}\) A positive voltage measured at the cold-probe (\(V_{\text{cold}}\)) with respect to the grounded hot-probe indicated that the deposited 3C-SiC films were n-type semiconductors with the carrier concentration (\(N_d\)) of approximately 10\(^{13}\) cm\(^{-3}\). Furthermore, as n-type 3C-SiC was employed in this work, the piezoresistors were aligned in [100] orientation to obtain the maximum magnitude of the effect.\(^{[33]}\) Figure 1(d) inset shows longitudinal and transverse 3C-SiC resistors aligned along [100] crystallographic direction.

The current-voltage (\(I-V\)) characteristic of the SiC piezoresistors was measured using Agilent B1500A, indicating that a good Ohmic contact was formed between Ni and n-type 3C-SiC. In addition, as the SiC was transferred onto glass, no current leakage was observed from the sensing layer to the substrate, as illustrated in Figure 2. The resistance of SiC was measured at different locations of the films (9 samples), showing a small deviation of less than 6%. This result indicates the uniformity of the carrier concentration throughout the highly doped film (See the supplementary document).

The bending beam method was applied to induce strain into the SiC piezoresistors, in which a SiC/glass cantilever was fixed, while the other free end was deflected downward by external forces.\(^{[34]}\) In addition, to avoid the effect of the boundary condition in the bending test, the clamped area was placed at least 2 mm far from the piezoresistors. The applied strain was estimated based on finite element analysis (FEA) using COMSOL Multiphysics. Accordingly, for the longitudinal resistance, nearly 100% of the strain applied to the top surface of the glass cantilever was transmitted to the SiC layer. On the other hand, for the transverse resistor, due to the small ratio of width to length of approximately 1:11, only 48% of applied strain was transmitted to the SiC layer (See supplementary material). This phenomenon of small strains induced into transverse resistors has also been reported in the literature, which is more profound for the case of nanowires with extremely small width to length ratio.\(^{[35]}\) Based on the FEA, when varying the applied force from 0 to 192 mN, the strains induced into longitudinal and transverse SiC piezoresistors were estimated to vary from 0 to 310 ppm, and from 0 to 140 ppm, respectively.

The response of the n-type 3C-SiC piezoresistors under applied strain was then investigated by monitoring the change in the output current under a constant applied voltage of 1 V, as shown in Figure 3(a). Accordingly, when increasing the tensile strain, the current passing through the SiC piezoresistors also increased. This characteristic in n-type 3C-SiC is different from p-type 3C-SiC as well as metals where the current decreases under tensile strain.\(^{[5,13]}\) The increase in the current in the n-type 3C-SiC also indicates the possibility to improve the mobility of n-type 3C-SiC on glass based devices by employing mechanical tensile strain. The change in the current also exhibits excellent repeatability after multiple bending cycles under an applied force of 192 mN (with high reproducibility of over 96% after 100 bending cycles). Figure 3(c),(d) plot the resistance change of the transverse resistor under applied loads, showing the opposite trend to that of the longitudinal resistor (i.e. resistance increased when increasing the tensile strain). Furthermore, it should be pointed out that under the same applied force to the SiC/Si cantilever, the change of the transverse resistance was relatively small in comparison to that of longitudinal resistance. This
result is considerable due to the fact that a smaller strain was induced into the transverse resistor as described above.

Figure 4 shows the resistance change of the n-type 3C-SiC on glass against applied strain to obtain the gauge factor of the material ($GF = \frac{\Delta R}{R_e}$). A linear relationship between the resistance change and tensile strain was observed in both longitudinal and transverse resistors. This linearity is a favorable property for strain sensing applications. Based on the results from Figure 4, the longitudinal and transverse gauge factors of n-type 3C-SiC were found to be $-8.6$ and $10.5$, respectively, which are at least five times larger than those of metals (See supplementary material).

Employing the theory of strained n-type cubic semiconductors, we qualitatively explain the piezoresistance of n-type 3C-SiC on glass based on the electron transfer phenomenon.[36,37] That is, under applied strain, the conduction bands of the n-type 3C-SiC are deformed, leading to the re-population of electrons, following the Boltzmann distribution. According to the first principle analysis of Nakamura et al. on a n-type 3C-SiC nanosheet,[38,39] under a tensile strain in [100] orientation, the energy valley in the [100] (i.e., the longitudinal valley) will shift up, while the energy valleys in [010] and [001] directions (i.e., the transverse valleys) will shift down. These shifts in the 6 energy valleys result in the re-population of free electrons, in which the electrons from the longitudinal direction (i.e., [100] orientation) will fill up the valley in transverse direction (i.e., [010] and [001] orientations). Moreover, since the transverse mobility of electron ($\mu_t$) is higher than that in the longitudinal direction ($\mu_l$), more electrons in the [010] and [001] direction leads to a higher effective mobility. This hypothesis is consistent with our results that the tensile strain increases the conductivity of the n-type 3C-SiC, indicating the enhancement of electron mobility with the assumption that the total number of electrons remains constant under strain for highly doped semiconductors.[40] It should also be pointed out that because the minima of the second lowest conduction band in 3C-SiC locates more than 0.3 eV above the bottom of the M valleys (the lowest minima), only the change in the lowest conduction sub-band contributes to the piezoresistive effect of n-type 3C-SiC.

Next, we utilize the theoretical model of Kanda developed for Si–another cubic single crystal[41] to numerically estimate the gauge factor in the highly doped 3C-SiC. For strain-free n-type
3C-SiC, the the number of electrons in each equivalent valley is given by:

\[ n_0 = N_c \times F_{1/2} \left( \frac{E_c - E_F}{k_BT} \right) \]

where \( N_c \) is the effective density of state in the conduction band; \( F_{1/2} \) is the Fermi-Dirac integral; \( E_c \) and \( E_F \) are the energy of each valley in the conduction band and the Fermi level; \( k_B \) is the Boltzmann constant; and \( T \) is the absolute temperature. Under applied strain \( \varepsilon \) in [100] orientation, the energy of the valley located in [100] axis will shift up while the energies in the other valleys in [010] and [001] axes will shift down an amount of \( \Delta E = \Xi_0 \varepsilon \). Here, \( \Xi_0 \) is an independent constant of the deformation energy. As a result, the change in electron concentration in each valley is:

\[ \Delta n = N_c \times \left\{ F_{1/2} \left( \frac{E_c + \Delta E - E_F}{k_BT} \right) - F_{1/2} \left( \frac{E_c - E_F}{k_BT} \right) \right\} \]

\[ \approx N_c \times \frac{\Delta E}{k_BT} \times \left( E_c - E_F \right) \]

where \( F_{-1/2} = \delta F_{1/2} / \varepsilon \). Consequently, based on the electron transfer phenomenon, the gauge factor of n-type 3C-SiC is given by:

\[ G = 1 + 2\nu + \frac{\Xi_0 (L - 1)}{3k_BT (2L + 1)} \times \left( 2 + \nu \right) \times \frac{F_{-1/2}}{F_{1/2}} \]

where \( L = \mu_t / \mu_l \) is the ratio of the transverse and longitudinal electron mobilities, and \( \nu \) and \( \nu \) are the Poisson ratio of 3C-SiC and the substrate, respectively. The Fermi-Dirac integral was calculated based on Chang-Elizabeth approximation while the deformation energy \( \Xi_0 = 6.3 \text{ eV} \) was obtained based on the theoretical work of Lambrecht et al. on full potential band calculation. Additionally, the mobility ratio \( L = 2.74 \) was obtained from the experimental work reported in ref. [47]. Substituting these values into Equation (3), the gauge factor in degeneracy doped 3C-SiC with carrier concentration of \( 5 \times 10^{19} \) to \( 10^{20} \text{ cm}^{-3} \) varies from \(-8.6\) to \(-14\) which is consistent with our experimentally measured value. This result indicates that the electron transfer mechanism can qualitatively and quantitatively explain the piezoresistive effect in the highly doped 3C-SiC.

3. Conclusion

In conclusion, this work reports the piezoresistive effect in highly doped 3C-SiC nanotube film transferred onto a 6-inch glass substrate, employing anodic bonding technique. Experimental data shows a negative gauge factor of approximately \(-8.6\) in the longitudinal [100] direction of the n-type 3C-SiC, which is in solid agreement with numerical analysis based on electronic transfer effect in cubic materials under stress. The large gauge factors, superior mechanical properties, excellent chemical inertness, and bio-compatibility of the transferred 3C-SiC on glass could open up promising opportunities for MEMS applications in harsh environments, as well as bio-sensing.

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

Acknowledgements

This work was partially funded by the linkage grants LP150100153 and LP160101553 from the Australian Research Council (ARC). This work was performed in part at the Queensland node of the Australian National Fabrication Facility, a company established under the National Collaborative Research Infrastructure Strategy to provide nano and micro-fabrication facilities for Australia’s researchers. H.P. Phan acknowledges research grants from the Australian Nanotechnology Network Overseas Travel Fellowship and Griffith University Postdoctoral Fellowship (GUPF).

Conflict of Interest

The authors declare no conflict of interest.

Keywords

MEMS, piezoresistance, silicon carbide, strain engineering, wafer bonding.

Received: April 6, 2018
Revised: June 30, 2018
Published online: July 22, 2018

[1] D. G. Senesky, B. Jamshidi, K. B. Cheng, A. P. Pisano, IEEE Sens. J. 2009, 9, 14721478.
[2] C. R. Eddy, D. K. Gaskill, Science 2009, 324, 1398.
[3] R. S. Okojie, A. A. Ned, A. D. Kurtz, W. N. Carr, IEEE Trans. Electron Devices 1998, 45, 785.
[4] Y. Wu, Y. M. Lin, A. A. Bol, K. A. Jenkins, F. Xia, D. B. Farmer, Y. Zhu, P. Avouris, Nature 2011, 472, 74.
[5] H.-P. Phan, Piezoresistive Effect of p-Type Single Crystalline SiC, Springer, Heidelberg, Germany 2017, ISBN: 978-3-319-55543-0.
[6] T. Dinh, H.-P. Phan, A. Qamar, P. Woodfield, N. T. Nguyen, D. V. Dao, JMEMS 2017, 26, 966.
[7] X. Li, S. Chen, P. Ying, F. Gao, Q. Liu, M. Shang, W. Yang, J. Mater. Chem. C 2016, 4, 6466.
[8] J. B. Wei, L. Wang, F. Gao, J. Zheng, B. Tang, W. Yang, J. Mater. Chem. C 2013, 1, 4514.
[9] F. Gao, J. Zheng, M. Wei, W. Yang, Chem. Commun. 2011, 47, 11993.
[10] J. Rabkowski, D. Pefitsis, H. P. Nee, IEEE Ind. Electron. Mag. 2012, 6, 17.
[11] D. Nakamura, I. Gunjishima, S. Yamaguchi, T. Itô, A. Okamoto, H. Kondo, S. Onda, K. Takatori, Nature 2004, 430, 1009.
[12] N. G. Wright, A. B. Horsfall, K. Vassilevski, Mater. Today 2008, 11, 16.
[13] H.-P. Phan, T. Dinh, T. Koizumi, T.-K. Nguyen, A. Qamar, T. Namazu, N.-T. Nguyen, D. V. Dao, Appl. Phys. Lett. 2016, 109, 123502.
[14] H.-P. Phan, T.-K. Nguyen, T. Dinh, G. Ina, A. R. Kermann, A. Qamar, J. Han, T. Namazu, R. Maeda, D. V. Dao, N.-T. Nguyen, Appl. Phys. Lett. 2017, 110, 14906.
[15] P. Hens, A. A. Zakharchew, T. Iakimov, M. Syvajärvi, R. Yakimova, Carbon 2014, 80, 823.
[16] S. A. Kukushkin, A. V. Osipov, N. A. Feoktistov, Phys. Solid State 2014, 56, 1507.
[17] L Wang, S. Dimitrijev, A. Fissel, G. Walker, J. Chai, L. Hold, A. Fernandes, N.-T. Nguyen, A. Iacopi, RSC Adv. 2016, 6, 16662.
[18] G. Ferro, Crit. Rev. Solid State Mater. Sci. 2015, 40, 56.
[19] S. E. Saddow, Silicon Carbide Biotechnology: A Biocompatible Semiconductor for Advanced Biomedical Devices and Applications, Elsevier, Amsterdam, Netherlands 2012.
[20] T. Yang, L. Zhang, X. Hou, J. Chen, K. C. Chou, Sci. Rep. 2016, 6, 24872.
[21] V. Naglieri, B. Gludovatz, A. P. Tomsia, R. O. Ritchie, Acta Mater. 2015, 98, 141.
[22] J. Bi, G. Wei, L. Wang, F. Gao, J. Zheng, B. Tang, W. Yang, J. Mater. Chem. C 2013, 1, 4514.
[23] C. Serre, A. Romano-Rodriguez, A. Perez-Rodriguez, J. R. Morante, L. Fonseca, M. C. Acero, R. Kogler, W. Skorupa, Sens. Actuators A: Phys. 1999, 74, 169.
[24] T. Suga, F. Mu, M. Fujino, Y. Takahashi, H. Nakazawa, K. Iguchi, Jpn. J. Appl. Phys. 2015, 54, 030214.
[25] C.-H. Wu, C. A. Zorman, M. Mehregany, IEEE Sens. J. 2006, 6, 316.
[26] K. N. Vinod, C. A. Zorman, A. A. Yasseeen, M. Mehregany, J. Electron. Mater. 1998, 27, L17.
[27] H.-P. Phan, H. H. Cheng, T. Dinh, B. Wood, T.-K. Nguyen, F. Mu, H. Kamble, R. Vadivelu, G. Walker, L. Hold, A. Iacopi, D. V. Dao, T. Suga, N.-T. Nguyen, ACS Appl. Mater. Inter. 2017, 9, 27365.
[28] H.-P. Phan, A. Qamar, D. V. Dao, T. Dinh, L. Wang, J. Han, P. Tanner, S. Dimitrijev, N.-T. Nguyen, RSC Adv. 2015, 5, 56377.
[29] D. V. Dao, H.-P. Phan, A. Qamar, T. Dinh, RSC Adv. 2016, 6, 21302.
[30] T.-K. Nguyen, H.-P. Phan, H. Kamble, R. Vadivelu, T. Dinh, A. Iacopi, G. Walker, L. Hold, N.-T. Nguyen, D. V. Dao, ACS Appl. Mater. Inter. 2017, 9, 41641.
[31] H.-P. Phan, D. V. Dao, N.-T. Nguyen, Micro and Nanomanufacturing Volume II, Chapter 9: Silicon Micro-/Nanomachining and Applications, Springer, Cham, Switzerland 2017, pp. 225.
[32] P. Tanner, L. Wang, S. Dimitrijev, J. Han, A. Iacopi, L. Hold, G. Walker, Sci. Adv. Mater. 2014, 6, 1542.
[33] K. Nakamura, IEEJ Trans. Sens. Micromach. 2016, 136, 465.
[34] A. Qamar, H.-P. Phan, J. Han, P. Tanner, T. Dinh, L. Wang, S. Dimitrijev, D. V. Dao, J. Mater. Chem. C 2015, 3, 8804.
[35] T. Toriyama, Y. Tanimoto, S. Sugiyama, Microelectromech. Syst. 2002, 11, 605.
[36] J. C. Doll, B. L. Pruitt, Piezoresistor Design and Applications, 1st edn., Springer, New York, USA 2013.
[37] G. L. Bir, G. E. Pikus, Symmetry and Strain-Induced Effects in Semiconductors, Wiley, New York, USA 1974.
[38] K. Nakamura, T. Toriyama, S. Sugiyama, in Proc. 27th Sensor Symp. Sensors, Micromach., Appl. Syst. 2010, 16.
[39] K. Nakamura, T. Toriyama, S. Sugiyama, Jpn. J. Appl. Phys. 2011, 50, 06GE05.
[40] Y. Sun, S. E. Thompson, T. Nishida, Strain Effect in Semiconductors: Theory and Device Applications, 1st edn., Springer, New York, USA 2009.
[41] Y. Kanda, Sens. Actuators A: Phys. 1991, 28, 83.
[42] Y. Kanda, Y. Kanda, IEEE Trans. Electron Devices 1982, 29, 64.
[43] T. Toriyama, S. Sugiyama, Appl. Phys. Lett. 2002, 81, 2797.
[44] H.-P. Phan, D. V. Dao, K. Nakamura, S. Dimitrijev, N.-T. Nguyen, J. Microelectromech. Syst. 2015, 24, 1663.
[45] C. Y. Chang, A. Izabelle, J. Appl. Phys. 1989, 65, 2162.
[46] W. R. L. Lambrecht, B. Segall, M. Methfessel, M. van Schilfessel, Phys. Rev. B 1991, 44, 3685.
[47] R. Kaplan, R. J. Wagner, H. J. Kim, R. F. Davis, Solid State Commun. 1985, 55, 67.