Effective optimization of surface passivation on porous silicon carbide using atomic layer deposited Al2O3

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1. Introduction

Boron (B) and nitrogen (N) co-doped SiC can provide yellow-orange light emission at room temperature. It has attracted significant attention as a candidate wavelength converter for SiC-based white light-emitting diodes (LEDs).\textsuperscript{1,2} To realize a pure white LED with a high color rendering index, an emission wavelength in the blue-green region is indispensable and porous SiC produced from bulk B–N co-doped SiC is a potentially attractive material. By using quantum confinement effect of B–N donor–acceptor-pair (DAP) recombination and surface defects in porous 6H-SiC, the emitting wavelength could be blue shifted to the blue-green spectra range.\textsuperscript{3–5} The optical and electronic properties of porous SiC are mainly determined by the morphological properties of porous structures, which have been investigated intensively using an anodic oxidation method.\textsuperscript{6,7} The morphology of porous SiC is usually determined by the experimental parameters such as the electrical current density, the composition of the etching solution and sample termination surface (Si or C) exposed to the etching solution.\textsuperscript{8}

Although porous SiC is an attractive wavelength converter, the luminescence of porous SiC is not very stable because of the large sensitive surface area. Even exposure to ambient air can cause degradation of PL due to the surface oxidation. Therefore, enhancement of the luminescence stabilization and improvement of luminescence intensity is necessary, and surface passivation of dangling bonds is an effective way. To date, still very little work has been reported to improve the passivation effect in porous SiC. Normally, there are two passivation strategies for semiconductor interface or surface: reduction of the non-radiative defect states (dangling bonds) by atomic H or thin dielectric films; significant reduction of the electron or hole concentration by negative-charge-dielectric films (introduce electric field).\textsuperscript{9} Al\textsubscript{2}O\textsubscript{3} films are known to provide outstanding effect of passivation on the surfaces of semiconductor, which can provide both of passivation strategies.\textsuperscript{9–14} For porous SiC, the important point to increase surface defects related photoluminescence is to reduce the non-radiative defect states (chemical passivation). Excess hydrogen introduced during the atomic layer deposition process is essential to the chemical passivation. The hydrogen atoms are usually bonded to the excess oxygen atoms and form –OH groups.\textsuperscript{12,15–18} It was observed that the passivation performance of ALD Al\textsubscript{2}O\textsubscript{3} can be activated by a post-annealing process.\textsuperscript{16} The surface passivation performance of Al\textsubscript{2}O\textsubscript{3} films was found to depend on several conditions, \textit{i.e.} the precursors,\textsuperscript{15,16} the deposition temperature\textsuperscript{16,22,23} and the post-deposition annealing temperature\textsuperscript{19,22,23} \textit{etc.} The result in our previous work on porous 6H-SiC without B doping has showed 250% enhancement of PL intensity.\textsuperscript{24} However, the passivation effect of ALD deposition processes, annealing conditions and the penetration depth on porous SiC haven't been investigated.

In this work, considering the importance of residual hydrogen content in the Al\textsubscript{2}O\textsubscript{3} for passivating porous SiC, we optimize the following parameters during the ALD Al\textsubscript{2}O\textsubscript{3} process: deposition temperature, Al\textsubscript{2}O\textsubscript{3} thickness and purging
time. The post-annealing temperature and duration are also investigated to avoid red shifting of the PL emission peak. The PL and XPS measurements were carried out to characterize the passivation effect in porous SiC. The optimization of passivation conditions on porous SiC is of great significance, as it provides a potential method to improve the luminescence efficiency in porous SiC.

2. Experimental

The porous 6H-SiC samples were prepared from single-crystalline 6H-SiC wafer (TakeneBlue, Beijing), doped with N (5.37 × 10^{18} \text{ cm}^{-3}) and B (0.95 × 10^{18} \text{ cm}^{-3}). Both sides of the substrate have been polished and have a Si terminating face (Si-face). A layer of 100 nm nickel was evaporated on the back side of the SiC samples to form an ohmic contact, followed by a layer of 10 nm titanium and 200 nm layer of gold. After that, the samples were mounted into a Teflon cell with a platinum counter electrode (cathode), as shown in Fig. 1. The anodic oxidation was processed under a pulsed current of 3 mA with a period of 50 ms and duty ratio of 50%. An aqueous solution with 5% HF was used for electrolytes. The fabrication process was carried under UV light illumination (mercury-xenon lamp; 4.5 W cm \(^{-2}\)) for 400 minutes. Under the pulsed current injection, HF electrolyte in the bath was circulated to reduce the adhesion of reactant CO\(_2\) bubbles. Finally, a porous layer with the thickness of 45 nm was obtained at the top surface. The color of the etched surface changed from transparent green to dark silver. For passivation optimization, porous samples a, b and c were fabricated under the same anodic etching conditions and further cleaved into small pieces (a\(_1\) to a\(_7\), b\(_1\) to b\(_{10}\) and c\(_1\) to c\(_4\)).

To investigate the passivation effect, the Al\(_2\)O\(_3\) films were deposited on the porous samples in an ALD reactor (Model R200, Piccusn, Finland), using trimethylaluminium (TMA:Al(CH\(_3\))\(_3\)) and H\(_2\)O gases as precursors for aluminum and oxygen, respectively. The precursors were alternately pumped into the reaction chamber in the ALD reactor by N\(_2\) carrier gas flow with the following sequence: 0.1 s TMA pulse, 3.0 s N\(_2\) purge, 0.1 s H\(_2\)O pulse, and 4.0 s N\(_2\) purge. One monolayer will be chemisorbed and retained on the surface of porous structures after one cycle. More details can be found in ref. 12 and 24. To study the passivation performance, 8 nm, 20 nm and 50 nm thick Al\(_2\)O\(_3\) were deposited at 160 °C for samples a\(_3\), a\(_6\) and a\(_7\) respectively. For comparison, samples a\(_2\), a\(_5\) and a\(_7\) were also covered with 8 nm, 20 nm and 50 nm thick Al\(_2\)O\(_3\) deposited at 200 °C. A post-deposition rapid thermal annealing (RTA) treatment of sample a\(_2\), a\(_3\), a\(_4\), a\(_5\), a\(_6\) and a\(_7\) were performed at 450 °C for 30 min in N\(_2\) (10 sccm) environment.

The second batch of porous SiC samples (sample b\(_2\) to b\(_{10}\)) covered with 20 nm thick of Al\(_2\)O\(_3\) were used to optimize the annealing temperature and duration. Finally, 20 nm thick Al\(_2\)O\(_3\) was deposited on the third batch of porous samples c\(_2\), c\(_3\) and c\(_4\) with 3.0, 20.0 and 30.0 s N\(_2\) purge time, respectively.

The morphology of the porous SiC before and after passivation was inspected by scanning electron microscopy (SEM). The crystal structure in porous SiC was recorded by X-ray diffraction (XRD). The passivation effect can be observed visually from the enhancement of PL intensity. To investigate the annealing effect on passivation, XPS was used to distinguish the elemental composition. The penetration depth of Al\(_2\)O\(_3\) films in the porous sample was analyzed by using a tunable beam of argon ions, then cycles of sputtering and measurement performed to measure the Al\(_2\)O\(_3\) composition as a function of sample depth.

3. Results and discussion

3.1 Morphology of the porous SiC covered with Al\(_2\)O\(_3\)

The cross-sectional SEM images of porous SiC without and with 20 nm thick Al\(_2\)O\(_3\) are shown in Fig. 2. The porous layer contains nano-triangular pores (marked by yellow triangle in Fig. 2(b)) with the size varying from 10–30 nm. The pore morphology varied across the depth of the sample from the triangular to the dotted line. This is attributed to the non-uniformity of the localized electrode potential and electrolyte transportation in deep pore structures. However, the pores structures of porous SiC in this work are more uniform than that fabricated by direct current and without UV illumination.\(^{24}\) The triangular porous line up in an orderly arrangement and then pile up layer by layer. The distance between the two layers is constant. In addition, the neighboring layers are in some cases connected by nearly vertically oriented pores (labeled with an ellipse in Fig. 2(b)). It has been demonstrated that the formed anodized porous layer is single-crystal 6H-SiC.\(^{25}\) Thus, the triangular pore can be used to identify the crystallographic orientation of the preferential etch planes and their variations with depth. Fig. 2(b) also shows very small structures (microcrystallites) between two adjacent pores which conform to the condition of quantum confinement with dimensions of 2–5 nm. However, for 6H-SiC (energy bandgap of 3.1 eV), the luminescence peak of quantum structure should be located in UV range, which will not be investigated in this work. After depositing 20 nm thick Al\(_2\)O\(_3\), the pores in the top layer are almost filled. The passivation boundary is not clearly observed in the SEM image, but it
appears that the bottom layer is not covered with Al$_2$O$_3$. The passivation depth will be further investigated by XPS measurement.

### 3.2 X-ray diffraction characterization of porous SiC

The crystal properties of original flat and porous samples have been characterized by XRD patterns using a Rigaku SmartLab® XRD system, with Cu Kα radiation from a Cu source ($\lambda = 1.5406$ Å). There are two obvious reflections from the crystal planes of 6H-SiC matrix, as shown in Fig. 3(a). The two main peaks are located at the angles of 35.60° and 75.38°, which corresponding to the planes (006) and (0012) of 6H-SiC, respectively. Details of the intensive XRD peaks located at 35.60° are presented in Fig. 3(b). It could be seen that two additional shoulder peaks (35.40° and 35.80°) appeared in the porous samples. These peaks can be attributed to the reflection from surface inclusions on triangular pores, such as dangling bonds and Si related oxide. After passivated by 20 nm Al$_2$O$_3$, the symmetric shoulder peaks became sharper due to the saturation of dangling bonds.

### 3.3 Al$_2$O$_3$ thickness optimization

To optimize the thickness, a layer of 8 nm, 20 nm and 50 nm thick Al$_2$O$_3$ was deposited on porous sample a$_2$, a$_3$ and a$_4$ at 160 °C, respectively. In addition to this, porous samples a$_5$, a$_6$ and a$_7$ were also covered with 8 nm, 20 nm and 50 nm thick Al$_2$O$_3$ deposited at 200 °C, and followed by a 30 min annealing at 450 °C. The passivation effect was characterized by PL measurements using a 375 nm laser as excitation source coupled into a microscope. After passivation, the PL intensity of porous samples a$_2$, a$_3$ and a$_4$ increased by a factor of 105%, 206% and 33% compared to reference sample a$_1$, respectively, as shown in Fig. 4(a). Fig. 4(b) shows that sample a$_6$ also has the largest PL enhancement of 178% among sample a$_5$, a$_6$ and a$_7$. However, the emitting peak of passivated porous samples is red shifted with 50 nm thick Al$_2$O$_3$. The cross-sectional SEM image in Fig. 2(d) shows that 20 nm of Al$_2$O$_3$ already fills the top layer of the porous sample. Excess oxygen atoms in 50 nm thick Al$_2$O$_3$...
may oxidize the porous surface and increase the oxygen-related defects during the annealing process, leading to the red shift. Therefore, it reveals that the effective passivation thickness is 20 nm for the porous SiC samples.

3.4 Impact of post-annealing conditions

The surface passivation quality of Al2O3 films grown by ALD was investigated after annealing under various conditions, i.e. annealing with different time and temperatures ranging from 300 °C to 500 °C. Based on the passivation results, all the samples were covered with 20 nm Al2O3 films deposited at 160 °C. Fig. 5 illustrates the passivation effect by PL measurement. Samples b2, b3, b4 and b5 were annealed for 5 min, 15 min, 30 min and 45 min at 450 °C, which resulted in an enhancement by a factor of 5.70, 5.57, 4.63 and 3.99, respectively, as shown in Fig. 5(a). Afterwards, the annealing temperature was also investigated for sample b6, b7, b8, b9 with 5 min at 300 °C, 350 °C, 400 °C, 450 °C and 500 °C, respectively. In Fig. 5(b), it can be seen that the enhancement for the passivated sample b6, b7, b8, b9 and b10 is 2.79, 5.40, 5.89, 5.70 and 3.03, respectively, compared to the reference sample b1. However, the emission peak of sample b6, b7, b8 and b10 is slightly red shift due to the oxidation effect with higher annealing temperature. The surface passivation performance was found to depend on both the annealing temperature and the duration. For higher annealing temperatures and longer annealing times the passivation performances degrades. It also can be deduced that the emission in porous 6H-SiC is mainly attributed to the surface defects.

3.5 Optimization of ALD purge time

From the cross-sectional SEM images in Fig. 2, it can be seen that the pore size is around 15–30 nm and only the pores in the top layer are entirely filled by Al2O3. To verify the ALD penetration depth and how it affects the PL, different purge time in the ALD process was studied. Porous samples c2, c3 and c4 were deposited with the purge time of 3.0 s, 20.0 s and 30.0 s, respectively. The PL spectra are shown in Fig. 6(a) where the original porous sample c1 (40 μm thick of porous layer) is used for reference. A passivation, the PL enhancement for sample c2, c3 and c4 are 3.43, 5.50 and 2.75, respectively. Thus, 30.0 s purge time is too long for the passivation due presumably to oxidation in the ALD react chamber.

The cross-sectional PL distribution was also characterized to inspect the penetration depth, as shown in Fig. 6(b). The excitation laser is guided into a microscope with 20× objective. The acquisition spot size is 1.14 μm, while the collected illumination signal is integration from 520 nm to 600 nm within the same porous depth. Thus the results in the cross-sectional PL are not comparable with that in the normal PL spectra in Fig. 6(a). The distribution curve for reference sample c1 shows that the emitted light intensity is uniform through the porous layer. It can be seen that the PL
intensity in sample c3 is higher than the other two samples within the top 6 μm layer. However, the maximum point in sample c3 and c4 is located about 6 μm and 16 μm beneath the surface respectively, which is attributed to the oxidation in the ALD chamber with much longer deposition time. Nevertheless, this trend is used to roughly judge the penetration depth of ALD deposition. A more accurate characterization for the optimized sample c3 will be shown by XPS characterization in the following section.

3.6 X-ray photoelectron spectroscopy characterization

XPS measurements were performed to investigate the passivation mechanisms and Al2O3 penetration depth in porous SiC. Fig. 7 shows Al 2p core-level XPS spectra in as-deposited and annealed porous SiC samples with Gaussian fitting. The pass energy was set to 50.0 eV for the measurements. The spectra were calibrated using C 1s at 284.8 eV as the internal reference to correct for charging. For as-deposited Al2O3 on porous SiC, two components can be observed at the binding energies of 76.04 eV and 76.9 eV, corresponding to Al–O bonds and Al–OH bonds, respectively. After annealing for 5 min, the component at 76.9 eV decreased, indicating the reduction of Al–OH bonds during post-deposition annealing, as shown in Fig. 7(b). It should be noted here that the residual Al–OH bonds existed during ALD deposition, which is easily broken and released within Al2O3 films. These released H atoms play an important role for the passivation performance in the porous layer. Such H atoms will diffuse and arrive at the Al2O3/SiC interface and thus neutralize the dangling bonds.17,23 This phenomenon is in consistent with that in XRD results.

In order to characterize the Al2O3 penetration depth, the porous sample c3 was analyzed by XPS with argon ion sputtering time, using ion beam energy of 3 keV. The acquisition spot size for the measurement is 400 μm. The results are shown in Fig. 8. Before sputtering, the intensity of Al 2p peak at 74.5 eV was the mixed signal of Al2O3 and absorbed CO2 on the surface. The peak was slightly shifted to 75.2 eV as the CO2 was sputtered off. However, the Al–O peak in Al 2p remains at the same binding energy location without discernible change with increasing sputtering time. The Al–O peak in O 1s gradually shifts to a higher binding energy with increasing sputtering time, and the intensity also decreases due to the Al2O3 thickness decreasing in the deeper porous layer. Based on the sputtering rate, both Al 2p and O 1s spectra could be used to qualitatively judge the passivation depth. The sputtering depth was measured to be 6.5 μm using a profiler. When zooming in the signal in Al 2p with a sputtering time of 45 min, the signal of Al–O bonds is still present. Thus the penetration depth for porous sample c3 is at least 6.5 μm. Simultaneously, the Si–C signal in C 1s and Si 2p is increased with the sputtering time. In addition, it is accounted for the contamination caused by the air exposure, which leads to bonding of the surface carbon atoms with the oxygen or water molecules from the air.

3.7 Time-resolved PL characterization

To analyze the passivation effect, time-resolved PL measurements were used to characterize the porous samples. The measurements were performed at room temperature. For the
excitation, a pulsed diode laser (pulse width \(\approx 44\) ps, 0.74 mW) working at 375 nm, controlled by a programmable laser driver (LDH-D-C-375 with PDL 820, PicoQuant GmbH), was focused onto the samples through a microscope. A burst mode excitation sequence was used to achieve a good signal-to-noise ratio in the measurement. In the burst mode, a pulse pattern was generated using the PDL 820 laser driver, consisting of 1 laser pulse followed by a waiting period of 62.5 ns, resulting in a total frequency of 16 MHz. The time-resolved PL results are presented in Fig. 9. The mechanisms of lifetime quenching are revealed by the lifetimes of the respective triple-exponential fitting, which includes non-radiative and radiative surface recombination and recombination in bulk SiC. The luminescence decay can be adequately described by triple-exponential fits, as shown in Table 1 and Fig. 9 with dotted lines. The average PL lifetime was calculated by using 

\[
\tau_{\text{ave}} = \frac{\sum A_i \tau_i^2}{\sum A_i \tau_i},
\]

<table>
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<th>Samples</th>
<th>(\tau_1) (ns)</th>
<th>(A_1)</th>
<th>(\tau_2) (ns)</th>
<th>(A_2)</th>
<th>(\tau_3) (ns)</th>
<th>(A_3)</th>
<th>(\tau_{\text{ave}}) (ns)</th>
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<tr>
<td>6H-Flat</td>
<td>11.93</td>
<td>0.061</td>
<td>2.87621</td>
<td>0.7283</td>
<td>0.55017</td>
<td>12.9097</td>
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<tr>
<td>(c_1)</td>
<td>6.52159</td>
<td>0.0185</td>
<td>1.80673</td>
<td>0.52743</td>
<td>0.34148</td>
<td>135.8705</td>
<td>0.3843</td>
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<tr>
<td>(c_2)</td>
<td>9.68234</td>
<td>0.01402</td>
<td>2.76457</td>
<td>0.48112</td>
<td>0.51831</td>
<td>28.18808</td>
<td>0.7815</td>
</tr>
<tr>
<td>(c_3)</td>
<td>8.27814</td>
<td>0.01635</td>
<td>2.50885</td>
<td>0.50592</td>
<td>0.46771</td>
<td>39.41452</td>
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<tr>
<td>(c_4)</td>
<td>7.91918</td>
<td>0.01458</td>
<td>2.38284</td>
<td>0.49494</td>
<td>0.45355</td>
<td>40.98371</td>
<td>0.6113</td>
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</table>
where $t_i$ and $A_i$ ($i = 1, 2, 3$) represent the lifetime and weight factor of each component. The average lifetime in bulk 6H-SiC and porous samples $c_1$, $c_2$, $c_3$ and $c_4$ is $1.89$ ns, $0.38$ ns, $0.78$ ns, $0.65$ ns and $0.61$ ns, respectively. Surface defects related non-radiative recombination quenches the PL lifetime in the porous samples. As expected the lifetime is increased to surface passivation, so the average lifetime in sample $c_2$ increased after passivation. It appears that the increase of ALD purge time results in a decrease of such radiative recombination channels. For longer purge times, the top layer of the porous sample might be oxidized in the ALD chamber thereby decreasing the passivation effect.

4. Conclusions

Porous B–N co-doped SiC samples were prepared by anodic oxidation of 6H-SiC substrate. The PL emission peak is blue shifted to around 520 nm due to surface defects and quantum confinement effect of B–N donor-acceptor-recombination in porous 6H-SiC. The XRD patterns show shoulder peaks of reflection from surface inclusions on the porous samples. Significant enhancement of the PL intensity was observed from porous SiC after deposition and annealing of ALD Al$_2$O$_3$ films. The thickness, annealing time, annealing temperature, and precursor purge time were optimized in term of strong PL. In order to study the passivation mechanism and penetration depth, XPS measurements were performed to check the chemical bonds in passivated porous SiC. It has been confirmed that the passivation effect is attributed to the OH bonds in Al$_2$O$_3$ films, which suppressed the non-radiative recombination after annealing. Moreover, the average carrier lifetime in Al$_2$O$_3$ passivated samples is longer than original porous SiC. However, there is a trade-off between the passivation and the oxidation effect during ALD deposition and annealing process. As a consequence, we draw the conclusions that the optimized passivation condition for porous SiC is 20 nm thickness of Al$_2$O$_3$, deposited at 160 °C and with a purge time of 20 s, followed by an annealing for 5 min at 350 °C. In addition, the red-shift phenomenon indicates that emission in porous SiC is mainly attributed surface defects. Such fundamental investigation provides valuable information to understand the emission mechanism and improve the emission efficiency in porous SiC based optical applications.

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Notes and references


