

Improved Performance and Reliability of N₂O-Grown Oxynitride on 6H-SiC

J. P. Xu, P. T. Lai, C. L. Chan, B. Li, and Y. C. Cheng

Abstract—This letter reports, for the first time, N₂O-grown oxides on both n-type and p-type 6H-SiC wafers. It is demonstrated that the N₂O-grown technique leads to not only greatly improved SiC/SiO₂ interface and oxide qualities, but also considerably enhanced device reliabilities as compared to N₂O-nitrided and conventional thermally-oxidized devices. These improvements are especially obvious for p-type SiC MOS device, indicating that N₂O oxidation could be a promising technique for fabricating enhancement-type n-channel SiC MOSFET's.

Index Terms—Hot-carrier stress, MOS devices, MOSFETs, nitridation, SiC.

I. INTRODUCTION

ONE OF the most important advantages of silicon carbide (SiC) over other wide-bandgap semiconductors is that it can be thermally oxidized to form SiO₂. However, it has been frequently reported that oxides grown on SiC have high interface-state density and a large amount of fixed oxide charges, especially for the SiO₂/p-SiC structure [1], [2]. Recently, NO and N₂O nitridation techniques have been used to improve the interface and oxide qualities [3], [4], and it is successful for NO-nitrided MOS devices. Unfortunately, it is found that N₂O nitridation deteriorates the electrical properties of devices. Interestingly in this work, an attractive method involving N₂O to effectively overcome the above problems is developed. For the first time, N₂O-grown technique used for Si MOSFET's is successfully applied to SiC MOS devices, and considerable improvements are obtained in not only SiO₂/SiC interface and oxide qualities, but also device reliability. Relevant results on N₂O-grown MOS capacitors are reported, suggesting that a high-quality and high-reliability SiO₂-SiC system could be realized by this N₂O-grown technology, even on p-type SiC substrate.

II. EXPERIMENTS

N- and p-type (0001) Si-face 6H-SiC wafers, manufactured by CREE Research, were used in this study. The SiC wafers had a 5- μ m epitaxial layer grown on heavily doped substrates. The doping level of the epitaxial layer of the two types of wafers was $4 \times 10^{15} \text{ cm}^{-3}$. The wafers were cleaned using the conventional RCA method followed by a 60-s dipping in 1% HF,

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and then loaded into a quartz furnace at 850 °C. After raising to an oxidation temperature of 1100 °C, a 360-min oxidation was carried out in pure N₂O ambient (denoted as N2OG sample). N- and p-type control samples were thermally oxidized in a wet O₂ ambient by bubbling oxygen through de-ionized water at 95 °C at the same temperature for 150 min (denoted as OX sample), because wet oxide is known to have better qualities than dry oxide [5], [6]. For the purpose of comparison, some n- and p-type OX samples were nitrided in pure N₂O ambient at 1100 °C for 1.5 h (denoted as N2ON sample). Then the three kinds of samples were annealed in N₂ at 1100 °C for 1 h. Finally, aluminum was thermally evaporated, and then patterned as gate electrode of MOS capacitors (area was $1.8 \times 10^{-4} \text{ cm}^2$), followed by a forming gas anneal at 410 °C for 30 min. High-field stress ($\pm 7 \text{ MV/cm}$), with the capacitor biased in accumulation, was used to examine device reliability. For comparing with the wet oxides, two other n-type wafers were oxidized in dry O₂ at the same temperature, but for a longer time of 3.5 h to get the same oxide thickness (denoted as DRY sample). One of them then went through the same N₂O nitridation as the wet samples. The SiO₂/SiC interface and oxide properties were characterized by flat-band voltage, oxide charge and their changes after stressing which were extracted from high-frequency (HF) capacitance-voltage (*C-V*) measurements at room temperature under dark condition. All measurements were carried out under a light-tight and electrically shielded condition.

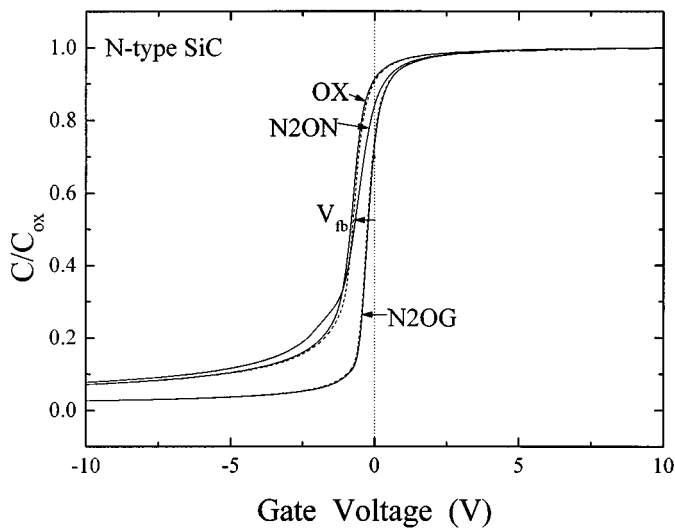
III. RESULTS AND DISCUSSIONS

Fig. 1 shows the typical HF *C-V* curves of all the samples under dark condition, swept in both directions, and no appreciable hysteresis occurs, implying little slow traps in the oxides. For n-type substrate, N2OG and N2ON samples exhibit slightly improved flat-band behavior relative to OX sample, with N2OG sample closest to the ideal *C-V* curve (flat-band voltages V_{fb} is almost 0 V). Obviously for p-type substrate, N₂O nitridation deteriorates the *C-V* properties of the device and a large difference between the two *C-V* curves for different sweep directions occurs, in consistent with the results in [3]. On the other hand, N₂O-grown sample exhibits an improved *C-V* behavior as compared to OX sample. Some parameters extracted from the HF-*C-V* curves are summarized in Table I. The oxide thickness t_{ox} is calculated from the HF accumulation capacitance or oxide capacitance C_{ox} . V_{fb} is determined from the flat-band capacitance C_{fb} [7]

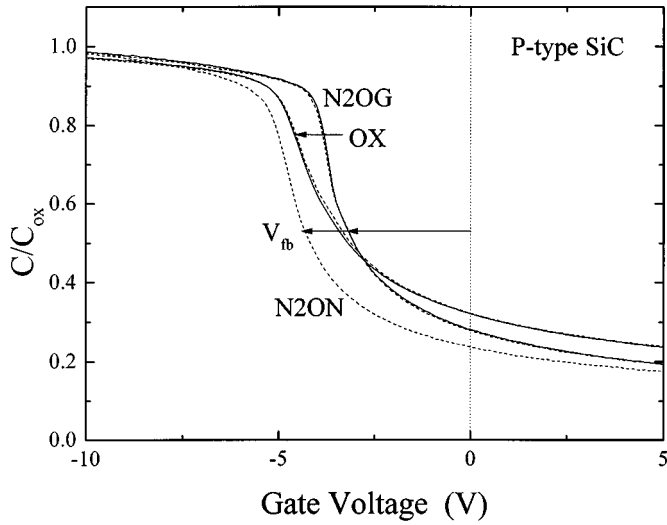
$$\frac{C_{fb}}{C_{ox}} = \left(1 + \frac{136 \sqrt{T/300}}{t_{ox} \sqrt{N}} \right)^{-1} \quad (1)$$

TABLE I
ELECTRICAL AND PHYSICAL PARAMETERS EXTRACTED FROM HF C-V CURVES

Sample	N-type SiC			P-type SiC		
	N2OG	N2ON	OX	N2OG	N2ON	OX
t_{ox} (Å)	206	217	210	203	213	200
C_{fb}/C_{ox}	0.52	0.53	0.52	0.51	0.53	0.51
V_{fb} (V)	-0.21	-0.65	-0.81	-3.05	-4.31	-3.40
$Q_{ox} \times 10^{11}$ cm ⁻²	2.5	5.9	8.2	5.4	16.7	9.4



(a)



(b)

Fig. 1. High-frequency C-V curve of all capacitors under dark condition, swept in two directions: depletion to accumulation (dashed line) and accumulation to depletion (solid line): (a) n-type sample and (b) p-type sample. Area of capacitor is 1.8×10^4 cm².

with N being acceptor or donor doping density and T the temperature in Kelvin. Oxide-charge density is calculated

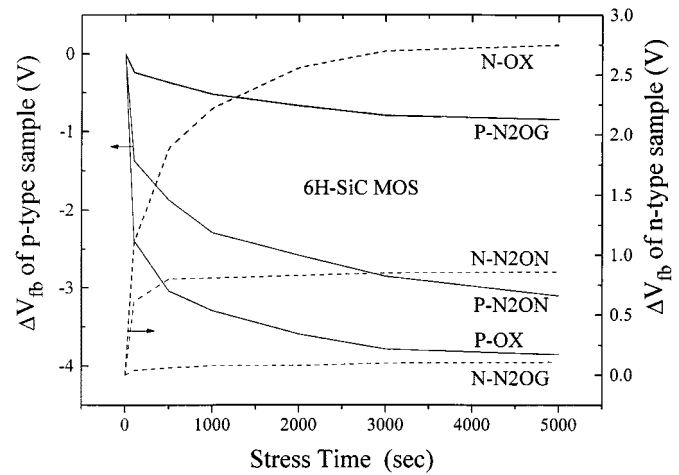


Fig. 2. Flat-band voltage shift of all samples under high-field stressing (± 7.0 MV/cm) at room temperature. P-means p-type sample while N-means n-type sample. Area of capacitor is 1.8×10^4 cm². Stress field = (gate voltage $-V_{fb}$)/ t_{ox} .

as $Q_{ox} = -C_{ox}(V_{fb} - \phi_{ms})/q$, where the work-function difference ϕ_{ms} between aluminum and 6H-SiC is calculated to be 0.031 V for n-type SiC and -2.53 V for p-type SiC. Unlike p-type N2ON sample, n-type N2ON sample also presents slightly improved C-V behavior, which is associated with pre-existing nitrogen at the SiO₂/SiC interface arising from the nitrogen-doped n-type substrate. Naturally, quality improvement of p-type N2OG sample can be deduced to come from nitrogen incorporation near the interface during N₂O oxidation, which is further confirmed by the stressing results below.

Presented in Fig. 2 are the flat-band voltage shifts of all the samples during a high-field stressing ($+7$ MV/cm for n-type samples and -7 MV/cm for p-type samples) at room temperature. Negative/positive ΔV_{fb} means generations of donor-type/acceptor-type interface states and positive/negative oxide charges. A much smaller ΔV_{fb} is obtained for both n-type N2OG and p-type N2OG samples, implying significantly suppressed generations of interface states and oxide traps in the two devices. In fact, the 0.1-V shift of the former after 5000 s is also much smaller than 0.4 V of DRY sample (not shown). Moreover, after N₂O nitridation, the shift of DRY sample further increases to 0.8 V, indicating a deterioration

of its interface [3]. From the maximum ΔV_{fb} at 5000-s stress time, increase of effective oxide charge is estimated to be $2.7/4.1 \times 10^{12} \text{ cm}^{-2}$ for n/p-OX samples, $7.5/29 \times 10^{11} \text{ cm}^{-2}$ for n/p-N2ON samples and $1.1/8.8 \times 10^{11} \text{ cm}^{-2}$ for n/p-N2OG samples. The greatly enhanced resistance of N2OG devices against the high-field stress should result from appreciable nitrogen incorporation at/near the SiO₂/SiC interface during oxide growth [8], [9] and thus formation of strong Si–N and O–N bonds [10], just like Si MOSFET's with N₂O-grown oxide as gate dielectric [9]–[12]. The two N2ON samples, especially n-N2ON sample, also exhibit smaller ΔV_{fb} than OX sample (although p-type sample has poor prestress HF C–V characteristics), supporting the above suggestion. In addition, during the initial 100-s stress time, ΔV_{fb} should mainly be linked to preexisting donor-/acceptor-type interface states and near-interface hole/electron traps, which are both neutral due to electron/hole occupations before the stress. From this, it can be proposed that N2OG oxide has much fewer donor-/acceptor-type interface states and oxide traps than OX oxide. This is in good agreement with the results in Table I. Therefore, it can be suggested that N₂O-grown oxide on both n- and p-type SiC substrates can lead to not only fewer interface states and oxide charges, but also hardened interface and near-interface oxide due to nitrogen pile-up near the interface.

IV. SUMMARY

N₂O-grown oxide on both n- and p-type 6H–SiC wafers was successfully fabricated for the first time. Improved SiC/SiO₂ interface and oxide qualities were demonstrated in N₂O-grown MOS devices, especially for p-type devices. Moreover, con-

siderably enhanced device reliability under high-field stress was observed with much smaller flat-band voltage shift than N₂O-nitrided and thermal-oxide devices. All these gains are attributed to significant nitrogen incorporation near the SiC/SiO₂ interface during N₂O oxidation. Therefore, N₂O oxidation technique could have great potential in making high-quality and high-reliability SiC MOSFET's, especially n-channel ones.

REFERENCES

- [1] S. T. Sheppard, J. A. Cooper, Jr., and M. R. Melloch, "Non-equilibrium characteristics of the gate-controlled diode in 6H–SiC," *J. Appl. Phys.*, vol. 75, pp. 3205–3207, 1994.
- [2] D. Alok, P. K. McLarty, and B. J. Baliga, "Electrical properties of thermal oxide grown using dry oxidation on p-type 6H–silicon carbide," *Appl. Phys. Lett.*, vol. 65, pp. 2177–2178, 1994.
- [3] S. Dimitrijević, H. F. Li, H. B. Harrison, and D. Sweatman, "Nitridation of silicon-dioxide films grown on 6H silicon carbide," *IEEE Electron Device Lett.*, vol. 18, pp. 175–177, May 1997.
- [4] H. F. Li, S. Dimitrijević, and H. B. Harrison, "Improved reliability of NO-nitrided SiO₂ grown on p-type 4H–SiC," *IEEE Electron Device Lett.*, vol. 19, pp. 279–281, Aug. 1998.
- [5] S. M. Tang *et al.*, *J. Electrochem. Soc.*, 1990, vol. 137, p. 221.
- [6] S. Zaima, K. Onoda, Y. Koide, and Y. Yasuda, *J. Appl. Phys.*, 1990, vol. 68, p. 6304.
- [7] D. K. Schroder, *Semiconductor Material and Device Characterization*, 2nd ed. New York: Wiley, 1998.
- [8] T. Hori and H. Iwasaki, *J. Appl. Phys.*, 1989, vol. 65, p. 629.
- [9] W. Ting, H. Hwang, J. Lee, and D. L. Kwong, "Composition and growth kinetics of ultra-thin SiO₂ films formed by oxidizing Si substrates in N₂O," *Appl. Phys. Lett.*, vol. 57, p. 2808, 1990.
- [10] T. Y. Chu *et al.*, "Study of the composition of thin dielectrics grown on Si in a pure N₂O ambient," *Appl. Phys. Lett.*, 1991.
- [11] W. Ting *et al.*, "MOS characteristics of ultrathin SiO₂ prepared by oxidizing Si in N₂O," *IEEE Electron Device Lett.*, vol. 12, p. 416, 1991.
- [12] G. Q. Lo, W. Ting, J. Ahn, and D. L. Kwong, "Improved performance and reliability of MOSFET's with ultra-thin gate oxides prepared by conventional furnace oxidation of Si in pure N₂O ambient," in *Proc. VLSI Tech. Symp. Dig.*, 1991, p. 43.