

Effects of Wet Oxidation/Anneal on Interface Properties of Thermally Oxidized SiO₂/SiC MOS System and MOSFET's

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Abstract— Effects of wet atmosphere during oxidation and anneal on thermally oxidized p-type and n-type MOS interface properties were systematically investigated for both 4H- and 6H-SiC. Deep interface states and fixed oxide charges were mainly discussed. The wet atmosphere was effective to reduce a negative flatband shift caused by deep donor-type interface states in p-type SiC MOS capacitors. Negative fixed charges, however, appeared near the interface during wet reoxidation anneal. In n-type SiC MOS capacitors, the flatband shift indicated a positive value when using wet atmosphere. The relation between interface properties and characteristics of n-channel planar 6H-SiC metal-oxide-semiconductor field effect transistors (MOSFET's) was also investigated. There was little relation between the interface properties of p-type MOS capacitors and the channel mobility of MOSFET's. The threshold voltage of MOSFET's processed by wet reoxidation anneal was higher than that of without reoxidation anneal. A clear relation between the threshold voltage and the channel mobility was observed in MOSFET's fabricated on the same substrate.

Index Terms—Deep interface states, fixed oxide charges, interface, MOS, MOSFET's, reoxidation anneal, silicon carbide, wet oxidation.

I. INTRODUCTION

SILICON CARBIDE (SiC) is a promising semiconductor for high-power, high-temperature, and high-frequency devices because of its superior properties such as wide bandgap, high breakdown field, high saturation electron drift velocity, and high thermal conductivity. For the aspect of device processes, SiC has an advantage of both n- and p-type conduction in a wide range by either *in situ* doping [1], [2] or selective ion implantation [3], [4]. One of the most important advantages of SiC over other wide bandgap semiconductors is that SiC can be thermally oxidized to form high-quality SiO₂ like Si: metal-oxide-semiconductor field effect transistors (MOSFET's) can be made on SiC.

Nowadays, SiC MOSFET's have been intensively investigated by many groups [5]–[8]. However, their excellent characteristics expected from SiC properties have not been

achieved yet, mainly due to inferior SiO₂/SiC interface properties. While rather good characteristics are obtained on the SiO₂/n-SiC interface [9], [10], the SiO₂/p-SiC interface has a large amount of fixed charges and large interface state density [11], [12]. In recent years, wet oxidation [13] or wet reoxidation anneal [14] at low temperatures was reported to reduce the interface state density of p-type MOS capacitors. They did little discuss, however, on the effects of oxidation or anneal ambient on deep states and fixed charges near the interface. In addition, there have been few reports on the effects of oxidation or anneal ambient on SiO₂/n-SiC interface properties.

In this paper, we report systematic investigation on the effects of wet oxidation and reoxidation anneal in wet O₂ at low temperatures on the interface properties (deep interface states and fixed charges) of SiO₂/p-SiC and SiO₂/n-SiC, for both 4H and 6H polytypes. We also investigated on the relation between the interface properties and characteristics of n-channel planar 6H-SiC MOSFET's.

II. EXPERIMENTS

MOS capacitors were fabricated on homo-epilayers grown on a (0001) Si-face of both 4H- and 6H-SiC. B-doped p-epilayers and N-doped n-epilayers were grown on highly doped p-type and n-type substrates, respectively. The epilayer thickness was 4 μm. The impurity concentrations of epilayers evaluated from Schottky characteristics were $2 \times 10^{16} \text{ cm}^{-3}$ for p-type and $2 \times 10^{15} \text{ cm}^{-3}$ for n-type. In this experiment, B-doped epilayers were used for p-type samples because the controllability of low doping level for B-doped epilayers is better than Al-doped epilayers in our CVD system. However, there is no essential difference in SiO₂/SiC interface properties between B-doped and Al-doped samples according to previous reports [15], [16].

Before thermal oxidation, the samples were cleaned by RCA cleaning followed by 0.5% HF dip. Thermal oxidation was carried out at 1150 °C in dry O₂ for 3 h or in wet O₂ (bubbling O₂ through deionized water at 95 °C) for 2 h followed by post-oxidation anneal at 1150 °C in Ar for 30 min. The oxide thicknesses determined from accumulation in high-frequency capacitance–voltage (*C*–*V*) curve were 350–400 Å for dry oxidation and 300–350 Å for wet oxidation. For some samples, reoxidation anneal [14] was carried out at 950 °C for 3 h in wet O₂ after Ar anneal, and in cooling off in Ar ambient, the

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ramp rate was 3 °C/min. The samples were pulled out at room temperature. As a gate electrode, Al was evaporated through a metal mask to form circular (300, 500 μm in diameter) patterns. After forming the gate electrode, post-metallization anneal (PMA) was performed in a forming gas ($\text{N}_2 : \text{H}_2 = 9 : 1$) at 400 °C for 30 min.

N-channel planar MOSFET's were fabricated on p-type 6H-SiC epilayers using a non self-aligned process. The source and drain regions were formed using multiple N^+ ion implantations with a total dose of $5 \times 10^{14} \text{ cm}^{-2}$. Post-implantation anneal was carried out at 1500 °C for 30 min in Ar. The gate oxidation was carried out in the same manner as the fabrication of MOS capacitors above mentioned. The channel length (L) was either 10 or 30 μm , and the channel width (W) was 200 μm . Ohmic contacts for source and drain regions were Al without an alloy process. After forming the Al gate electrode, PMA was performed.

The SiO_2/SiC interface properties were characterized by high-frequency $C-V$ measurements of MOS capacitors at room temperature under the dark condition and after light illumination at deep depletion. The high-frequency $C-V$ characteristics were measured with a probing frequency of 100 kHz and a bias sweep rate of 0.1 V/s. The total deep interface states and fixed charges were estimated from a ledge feature caused by deep interface states and a flatband voltage shift. The SiO_2/SiC interface properties were also characterized by MOSFET parameters such as the channel mobility and threshold voltage at room temperature.

III. RESULTS AND DISCUSSION

A. $\text{SiO}_2/p\text{-SiC}$ Interface Properties

Fig. 1 shows typical high-frequency $C-V$ curves under the dark condition for p-type 4H-SiC MOS capacitors fabricated by different oxidation processes. The measured curves show deep depletion because of the extremely low intrinsic carrier concentration due to the wide bandgap. Although a large negative flatband shift (ΔV_{fb}) was observed for the dry oxidation samples, wet oxidation can reduce ΔV_{fb} . Furthermore, the wet reoxidation anneal can reduce ΔV_{fb} to less than -1.0 V. The reoxidation anneal was also useful even after dry oxidation to reduce ΔV_{fb} to the same value as after wet oxidation. This indicates that the interface properties of MOS capacitors processed by wet reoxidation anneal at a low temperature do not depend on the oxidation gas.

The large negative ΔV_{fb} in p-type MOS capacitors is mainly caused by holes trapped at deep donor-type interface states, and not by fixed charges [17]. If negative fixed charges exist near the interface, however, the flatband shift can be reduced. Deep interface states and fixed charges were tried to be separated for MOS capacitors processed by wet oxidation and wet reoxidation anneal as well as a dry-oxidized MOS capacitor using high-frequency $C-V$ characteristics after light illumination.

Fig. 2 shows a high-frequency $C-V$ curve for a dry-oxidized p-type 4H-SiC MOS capacitor measured after light illumination at deep depletion. A tungsten lamp with a power

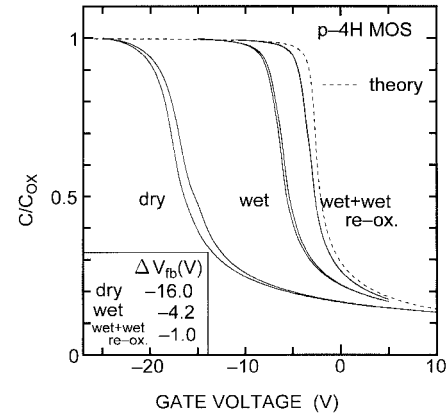


Fig. 1. High-frequency $C-V$ curves of p-type 4H-SiC MOS capacitors measured under the dark condition. The dotted line denotes a theoretical curve for deep depletion.

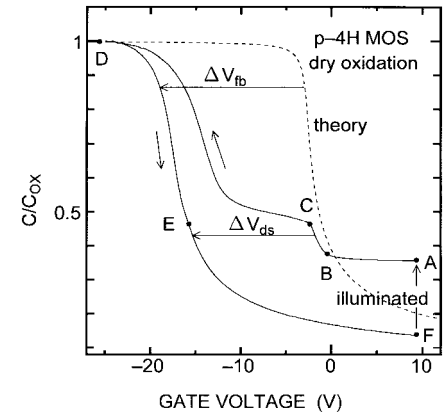


Fig. 2. High-frequency $C-V$ curve of dry-oxidized p-type 4H-SiC MOS capacitor measured after light illumination at deep depletion.

of 120 W was used as the light source for illumination. The MOS capacitor was illuminated for more than 1 min to form an inversion layer at a bias of 10 V. After the illumination was turned off, the bias was swept in the sequence of A→B→C→D→E→F. When the inversion layer is formed by the illumination, the deep interface states capturing holes start to capture electrons and change the charge state to neutral, since the deep interface states are donor-type. At point C, the $C-V$ curve shows a so-called “interface state ledge” [18] that appears when the deep states exist at the interface. At point D, holes accumulate near the interface and all deep interface states capture these holes. Once holes are captured at the deep interface states, they can not be thermally emitted to the valence band at room temperature. So, in the region of D→E→F, the charge state of the deep interface states is positive. The voltage shift between the region of B→C and that of E→F is caused by the difference of the charge states of the deep interface states. From this voltage shift (ΔV_{ds}), the total deep interface state density per unit area (N_{ds}) is obtained by

$$N_{\text{ds}} = \frac{C_{\text{ox}} |\Delta V_{\text{ds}}|}{q} \quad (1)$$

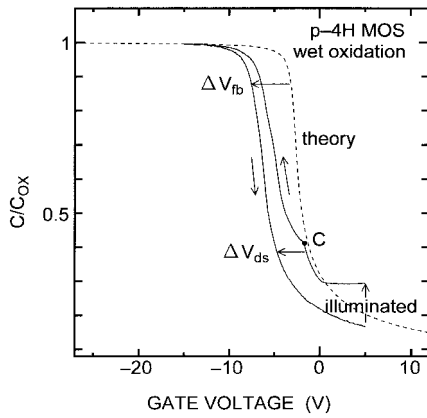


Fig. 3. High-frequency C - V curve of wet-oxidized p-type 4H-SiC MOS capacitor measured after light illumination.

Here, C_{ox} is the capacitance of the gate oxide and q the electronic charge. Fig. 2 indicates a ΔV_{ds} of -13.5 V, and the total deep interface state density N_{ds} is estimated to be $6.0 \times 10^{12} \text{ cm}^{-2}$ by (1). The flatband shift from the theoretical curve is caused by both holes trapped at the deep interface states and fixed charges. The subtraction of ΔV_{ds} from ΔV_{fb} gives the voltage shift caused by only the fixed charges. Then the fixed charge density is given by

$$Q_{fc} = \frac{C_{ox}(|\Delta V_{fb}| - |\Delta V_{ds}|)}{q} \quad (2)$$

In this dry-oxidized sample, the flatband shift was $\Delta V_{fb} = -16.0$ V. From these results, the fixed charge density Q_{fc} was estimated to be $1.1 \times 10^{12} \text{ cm}^{-2}$ by (2). Q_{fc} may include not only fixed charges but also other charges such as mobile ions and charged border traps besides deep interface states.

The small ΔV_{fb} of a wet-oxidized MOS capacitor in Fig. 1 could be realized by increase of negative fixed charges. Fig. 3 shows a high-frequency C - V curve of a wet-oxidized p-type 4H-SiC MOS capacitor. The point C indicating an "interface state ledge" is located near the negative side of the theoretical curve. If negative fixed charges exist near the interface, the ledge can be located at the positive side of the theoretical curve. Therefore, the increase of negative fixed charges was negligible or a few during the wet oxidation compared with a large reduction of N_{ds} . In this sample, ΔV_{fb} of -4.2 V and ΔV_{ds} of -3.2 V lead to N_{ds} of $1.9 \times 10^{12} \text{ cm}^{-2}$ and the significant reduction of N_{ds} contributes to the decrease of the large negative ΔV_{fb} , while the small reduction of Q_{fc} observed in the wet-oxidized sample compared with the dry-oxidized sample also contributes. The variation of fixed charges among samples might be caused by impurity contamination during Al evaporation. The same measurements were performed for p-type 6H-SiC using both dry- and wet-oxidized samples, and the results were similar to those of 4H-SiC.

Fig. 4 shows high-frequency C - V curves after light illumination for p-type MOS capacitors processed by wet oxidation with wet reoxidation anneal. The results for the sample processed by dry oxidation with wet reoxidation anneal were similar. A ledge (point C) was observed for 4H-SiC as in Fig. 4(a), which is located at the positive bias compared with the theoretical curve. This results indicate the appearance

TABLE I
INTERFACE PROPERTIES OF $\text{SiO}_2/\text{p-SiC}$ WITH DIFFERENT OXIDATION AND ANNEAL AMBIENT. EFFECTIVE FIXED CHARGES (Q_{eff}) WERE CALCULATED FROM THE FLATBAND VOLTAGE SHIFT (ΔV_{fb}).

oxidation / anneal	ΔV_{fb} (V)	Q_{eff} ($\times 10^{11} \text{ cm}^{-2}$)	N_{ds} ($\times 10^{11} \text{ cm}^{-2}$)	Q_{fc} ($\times 10^{11} \text{ cm}^{-2}$)
dry/Ar	-16.0	71	60	11
wet/Ar	-4.2	25	19	6
dry/wet re-ox.	-1.1	6	18	-12
wet/wet re-ox.	-1.0	7	20	-13

(a) p-4H MOS

oxidation / anneal	ΔV_{fb} (V)	Q_{eff} ($\times 10^{11} \text{ cm}^{-2}$)	N_{ds} ($\times 10^{11} \text{ cm}^{-2}$)	Q_{fc} ($\times 10^{11} \text{ cm}^{-2}$)
dry/Ar	-15.1	72	55	17
wet/Ar	-2.9	18	14	4
dry/wet re-ox.	-0.6	6	-	-
wet/wet re-ox.	-0.7	6	-	-

-: Difficult to evaluate exact values.

(b) p-6H MOS

of negative charges near/at the interface by wet reoxidation anneal. From the flatband shift and the hysteresis of the ledge feature, the negative fixed charges and total deep state density were estimated to be $-1.4 \times 10^{12} \text{ cm}^{-2}$ and $2.0 \times 10^{12} \text{ cm}^{-2}$, respectively. Deep acceptor-type interface states appear, and in this case, negative fixed charges are caused by both "real" negative fixed charges and electrons captured at the deep acceptor-type interface states. Thus, the small flatband shift in wet reoxidation annealed 4H-SiC is considered to be caused by cancel out of both positive charges by trapped holes and the negative charges, though it is difficult to separate these two effects.

For wet reoxidation annealed p-type 6H-SiC, the ledge feature is not clear compared with 4H-SiC as in Fig. 4(a), but, it is actually observed at a bias of -1 V as in Fig. 4(b). So, there might exist negative charges near/at the interface of p-type 6H-SiC.

The interface properties of p-type MOS capacitors for 4H- and 6H-SiC fabricated by dry/wet oxidation and with/without wet reoxidation anneal at a low temperature are summarized in Table I. As mentioned above, the wet oxidation is effective to improve the interface properties of p-type MOS capacitors. The improvement of interface properties by wet oxidation may be attributed to the elimination of unwanted carbon compounds. Thus, the investigation of carbon at the interface is necessary to clarify the effect of wet oxidation. Another effect of wet oxidation might be the passivation of dangling bonds at the interface by hydrogen. The wet reoxidation anneal at a low temperature (950 °C) can reduce the flatband shift. However, negative fixed charges appear in the oxide near the interface. The origin of negative fixed charges might be OH^- , which can not diffuse out from the interface at 950 °C, but it is not fully understood yet.

B. $\text{SiO}_2/\text{n-SiC}$ Interface Properties

Fig. 5 shows typical high-frequency C - V curves of n-type 4H-SiC MOS capacitors under the dark condition. The wet-oxidized sample has more positive flatband shift than the

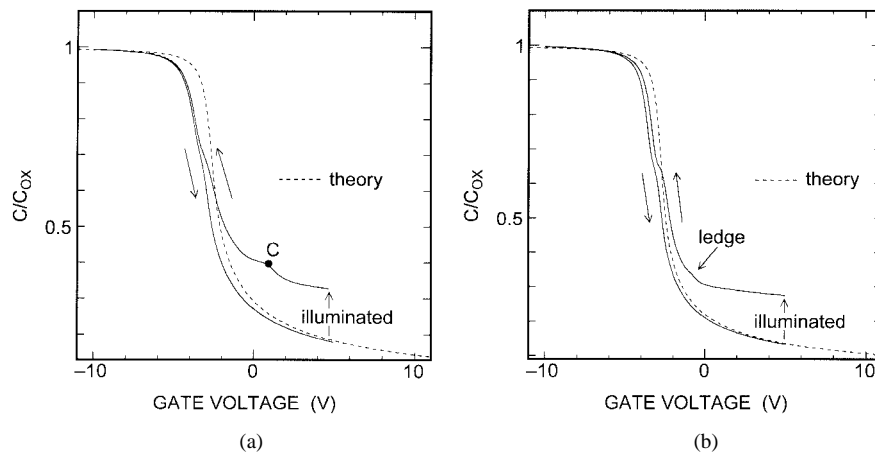


Fig. 4. High-frequency $C-V$ curve of wet reoxidation annealed p-type MOS capacitors measured after light illumination: (a) 4H-SiC MOS and (b) 6H-SiC MOS.

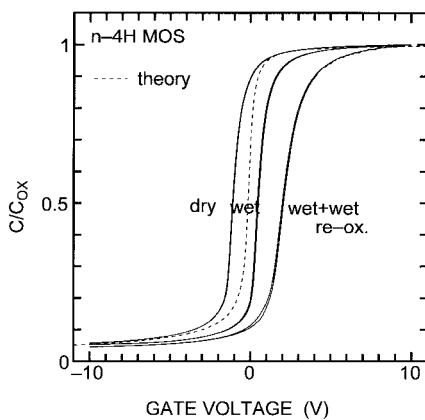


Fig. 5. High-frequency $C-V$ curves of n-type 4H-SiC MOS capacitors under the dark condition.

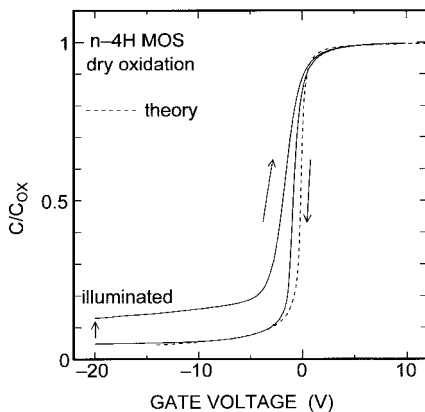


Fig. 6. High-frequency $C-V$ curve of dry-oxidized n-type 4H-SiC MOS capacitor after light illumination.

dry-oxidized sample. Still more, the flatband shift of the wet reoxidation annealed sample is more positive than that of samples without wet reoxidation anneal. When wet reoxidation anneal is done, the flatband shift takes almost the same value regardless of the oxidation ambient. The tendency is similar to that of p-type MOS capacitors.

Figs. 6 and 7 show high-frequency $C-V$ curves after light illumination for n-type 4H-SiC MOS capacitors processed by

dry and wet oxidation, respectively. For the dry-oxidized n-type sample, the ledge feature can not be seen as in Fig. 6, which means that there are few deep states at the $\text{SiO}_2/\text{n-SiC}$ interface. So, the flatband shift to the negative side for the dry-oxidized sample is mainly caused by real positive fixed charges. For the wet-oxidized n-type sample, on the other hand, the ledge feature can be seen at a bias of about -4 V as in Fig. 7, which means that deep acceptor-type interface states are formed by wet oxidation. It is difficult, however, to evaluate the deep interface states quantitatively because of dull ledge shape. Thus, only the flatband shift and effective fixed charges of both 4H- and 6H-SiC n-type MOS capacitors are given in Table II. These deep acceptor-type interface states capture electrons, and then, the charge state becomes negative. They may become one of the causes of the small positive flatband shift as in Fig. 5. Another cause of the positive flatband shift is negative fixed charges. For wet-oxidized p-type MOS capacitors, the reduction of positive fixed charges of $\text{mid-}10^{11} \text{ cm}^{-2}$ was obtained as in Fig. 3. If this reduction is caused by the increase of negative fixed charges, it is possible to increase negative fixed charges in n-type MOS capacitors by wet oxidation. Similar measurements were done for wet reoxidation annealed n-type samples, and the ledge feature could also be seen (not shown). Thus, the positive flatband shift shown in Fig. 5 is caused by both deep acceptor-type interface states and negative fixed charges. The same measurements were performed for n-type 6H-SiC processed by dry/wet oxidation and with/without wet reoxidation anneal, and the results were similar to those of 4H-SiC.

C. Relation Between Interface Properties and Characteristics of n-channel 6H-SiC MOSFET's

To investigate the relation between interface properties and characteristics of MOSFET's, 6H-SiC MOSFET's were fabricated using dry/wet oxidation and with/without wet reoxidation anneal. After all processes for fabrication of MOSFET's were done, the interface properties of MOS capacitors on the same chip of MOSFET's were confirmed to be the same as mentioned in Section III-A. The bias of both source and substrate was zero in the measurement of current-voltage characteristics.

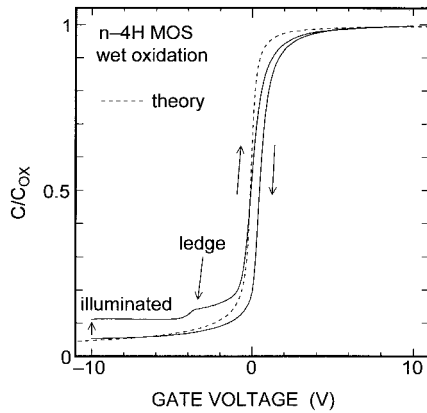


Fig. 7. High-frequency $C-V$ curve of wet-oxidized n-type 4H-SiC MOS capacitor after light illumination.

TABLE II
FLATBAND VOLTAGE SHIFT (ΔV_{fb}) AND EFFECTIVE FIXED CHARGES (Q_{eff}) OF n-TYPE MOS CAPACITORS WITH DIFFERENT OXIDATION AND ANNEAL AMBIENT

oxidation / anneal	ΔV_{fb} (V)	Q_{eff} ($\times 10^{11} \text{cm}^{-2}$)	ΔV_{fb} (V)	Q_{eff} ($\times 10^{11} \text{cm}^{-2}$)
dry/Ar	-0.9	5	-1.6	9
wet/Ar	0.7	-4	0.3	-2
dry/wet re-ox.	2.6	-17	1.4	-9
wet/wet re-ox.	2.3	-15	1.5	-10

(a) n-4H MOS (b) n-6H MOS

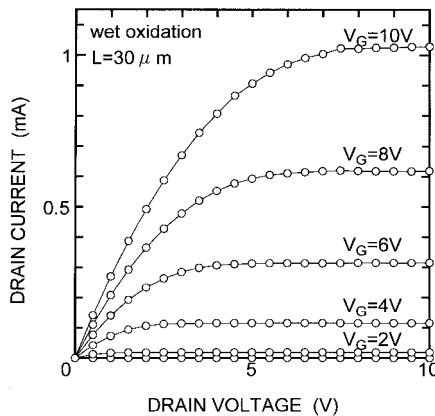


Fig. 8. Typical I_D-V_D characteristics of n-channel 6H-SiC MOSFET's using wet oxidation.

Typical drain current (I_D)-voltage (V_D) characteristics of 6H-SiC MOSFET's for different gate voltage (V_G)'s are shown in Fig. 8, which have good saturation and enhancement-mode characteristics. Other MOSFET's also showed good characteristics similar to Fig. 8. The channel mobility (μ_c) was determined from the linear region at low drain voltage in Fig. 8. Fig. 9 shows the channel mobility as a function of ($V_G - V_T$), where V_T is the threshold voltage. A maximum μ_c of $53.0 \text{ cm}^2/\text{Vs}$ was obtained for dry-oxidized and wet reoxidation annealed 6H-SiC MOSFET's, which is the best data at present in our experiments.

Fig. 10 shows typical $I_D^{1/2}-V_G$ characteristics of MOSFET's in the saturation region of $V_D = 10 \text{ V}$. Threshold voltage was determined by extrapolating the curve to zero

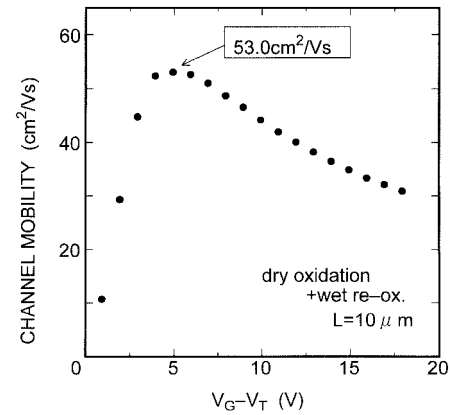


Fig. 9. Channel mobility as the function of ($V_G - V_T$). The maximum channel mobility of $53.0 \text{ cm}^2/\text{Vs}$ is obtained for dry-oxidized and wet reoxidation annealed 6H-SiC MOSFET's.

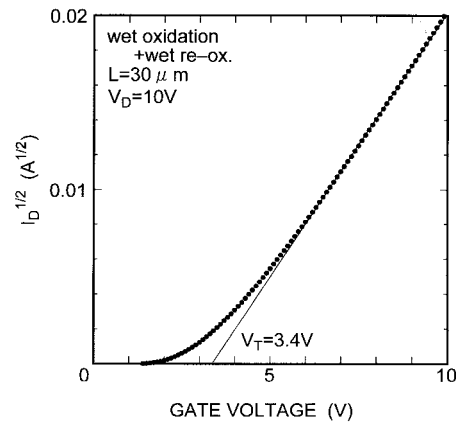


Fig. 10. Typical $I_D^{1/2}-V_G$ characteristics of n-channel 6H-SiC MOSFET's.

drain current, as illustrated in Fig. 10.

In the case of the existence of fixed oxide charges, the threshold voltage is given by

$$V_T = V_T(\text{theory}) - \frac{qQ_{nfc}}{C_{ox}}. \quad (3)$$

Here, $V_T(\text{theory})$ is the theoretical threshold voltage, and Q_{nfc} the net fixed charge density, which means the sum of both positive and negative fixed charges. The MOSFET parameters such as μ_c and V_T as well as Q_{nfc} for various interface properties are summarized in Table III: average of some (15–20) samples arbitrarily selected from one chip. From Table III, the followings are notified. First, the large negative flatband shift in MOS capacitors mainly caused by deep donor-type interface states has no relation with the threshold voltage of MOSFET's, because deep donor-type interface states can capture electrons in the inversion layer to be neutral. Second, deep donor-type interface states do not affect the channel mobility, which can be explained in the same reason above mentioned. Third, the threshold voltage of wet reoxidation annealed MOSFET's was higher than that of without reoxidation anneal. This indicates that negative charges appear near the interface by wet reoxidation anneal, that are the same results of the appearance of negative charges in MOS capacitors described in Section III-A.

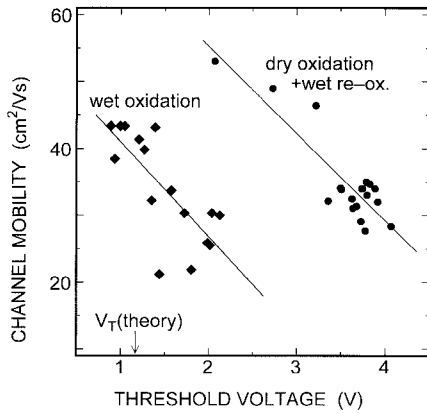


Fig. 11. Relation between channel mobility and threshold voltage on MOSFET's fabricated on the same substrate.

TABLE III
AVERAGE VALUES OF μ_c , V_T , AND Q_{nfc} FOR 6H-SiC
MOSFET'S PROCESSED BY VARIOUS OXIDATION CONDITIONS

oxidation / anneal	μ_c (cm ² /Vs)	V_T (V)	$V_T(\text{theory})^*$ (V)	Q_{nfc} ($\times 10^{11}$ cm ⁻²)
dry/Ar	34.1	1.4	1.66	1
wet/Ar	34.5	1.4	1.17	-2
dry/wet re-ox.	35.2	3.5	1.16	-13
wet/wet re-ox.	35.6	3.7	1.02	-17

*:Variation of $V_T(\text{theory})$ due to different oxide thickness.

Then, what does affect the channel mobility of SiC MOSFET's? It seems that one of the hints to answer this question is in Fig. 11. The MOSFET's fabricated on the same substrate have the tendency that the channel mobility becomes high if the measured threshold voltage becomes close to the theoretical value. In other words, it can be said that net negative charges near/at the interface cause to suppress the channel mobility of SiC MOSFET's, in spite of the difference in oxidation/anneal conditions in Fig. 11. The channel mobility is expected to be improved further, if wet reoxidation anneal condition is optimized and net negative charges near/at the interface can be reduced. Thus, the optimization of wet reoxidation anneal condition is necessary to achieve a higher channel mobility. Other causes such as shallow interface states near the conduction band edge, roughness at the interface and crystallinity of epilayers might be considered.

IV. CONCLUSIONS

The effects of wet oxidation/anneal on the interface properties of p-type and n-type MOS capacitors were systematically investigated for both 4H- and 6H-SiC. For the p-type MOS capacitors, wet oxidation was effective to reduce the deep donor-type interface states resulting in more small negative flatband shift than dry oxidation. The flatband shift of less than -1 V was obtained using wet reoxidation anneal. The negative fixed charges, however, appeared near the interface. For n-type MOS capacitors, there were few deep interface states on dry-oxidized samples. The flatband shift of wet-oxidized n-type samples indicated positive Q_{eff} due to the appearance of deep acceptor-type interface states and negative fixed charges, and these effects were emphasized by wet reoxidation anneal.

The relation between the interface properties and characteristics of n-channel planar 6H-SiC MOSFET's were also investigated. The large negative flatband shift caused by deep donor-type interface states has no relation with the threshold voltage and the channel mobility of MOSFET's. On the contrary, there was a clear relation between the threshold voltage and the channel mobility in MOSFET's fabricated on the same substrate. The threshold voltage of wet reoxidation annealed MOSFET's was higher than that of without reoxidation anneal due to the negative fixed charges near the interface.

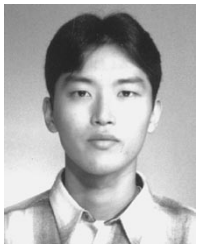
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