Surface Passivation of Crystalline Silicon by Combination of Amorphous Silicon Deposition with High-Pressure H₂O Vapor Heat Treatment

Toshiyuki Sameshima*, Tomokazu Nagao, Masahiko Hasumi, Asuka Shuku¹, Eiji Takahashi¹, and Yasunori Andoh¹

Tokyo University of Agriculture and Technology, Koganei, Tokyo 184-8588, Japan ¹NISSIN Electric Co., Ltd., Kyoto 615-8686, Japan

Received July 20, 2011; revised November 20, 2011; accepted November 24, 2011; published online March 21, 2012

A high minority carrier effective lifetime τ_{eff} of crystalline silicon was achieved by hydrogenated amorphous silicon (a-Si:H) films formed by a combination of plasma-enhanced chemical vapor deposition at 150 °C with high-pressure H₂O vapor heat treatment. τ_{eff} was 1.6 x 10⁻⁴, 3.0 x 10⁻⁴, and 1.15 x 10⁻³ s for n-type silicon substrates coated with 3-, 10-, and 50-nm-thick a-Si:H films treated with 1.0 x 10⁶ Pa H₂O vapor heat treatment between 180 and 300 °C for 1 h. Light-induced passivation enhancement was demonstrated when 620-nm light was illuminated at the 50-nm-thick a-Si:H surface. τ_{eff} increased from 8.5 x 10⁻⁴ to 1.15 x 10⁻³ s probably caused by field effect passivation induced by hole trapping at the SiO_x formed by H₂O vapor heat treatment for 1 h. On the other hand, τ_{eff} was further increased to 1.2 x 10⁻³ s by 1.0 x 10⁶ Pa H₂O vapor heat treatment at 300 °C for 3 h for the sample formed with the 50-nm-thick a-Si:H film. However, no increase in τ_{eff} was observed by light illumination at the a-Si:H surface, probably because the SiO_x clusters became stable and had no hole trapping property. (© 2012 The Japan Society of Applied Physics

1. Introduction

Hydrogenated amorphous silicon (a-Si:H) has been widely used to passivate the surface of crystalline silicon.¹⁻⁴⁾ The surface recombination velocity of photo-induced minority carriers is effectively decreased and the minority carrier effective lifetime τ_{eff} is increased by the formation of a-Si:H layers by plasma enhanced chemical vapor deposition (PECVD) because of dangling bond termination by hydrogen atoms included in a-Si:H layers at the crystalline silicon surface. a-Si:H passivation has been applied to the fabrication of solar cells with a high conversion efficiency such as hetero junction with intrinsic type (HIT) silicon solar cells.⁵⁾ However, we believe that it is not easy to directly form high quality a-Si:H and achieve a low recombination velocity because high energy ions and radicals in plasma can cause serious defects increasing the recombination velocity at the silicon surface.⁶⁾ Post annealing will be practical to improve silicon surface passivation with a-Si:H.

In this paper, we report a high τ_{eff} of crystalline silicon substrates achieved by a-Si:H films formed by PECVD at 150 °C followed by post high-pressure H₂O vapor heat treatment, which has been developed to reduce the density of defects at the SiO₂/Si interface and grain boundaries of polycrystalline silicon.^{7,8} Moreover, we report a lightinduced passivation phenomena. τ_{eff} increased by 30% when the a-Si:H surface was illuminated by light.

2. Experimental Procedure

 $30-\Omega$ cm n-type silicon substrates with a thickness of $520 \,\mu$ m and a crystalline orientation of (100) were prepared. The top and rear surfaces were coated with 100-nm-thick thermally grown SiO₂ layers formed at $1100 \,^{\circ}$ C in dry oxygen atmosphere. The SiO₂ layer at the top surface was then removed using 5% diluted hydrofluoric acid. a-Si:H films with average thicknesses of 3, 10, and 50 nm were deposited on the top surface.⁹⁾ The average film thickness was estimated by deposition duration with the deposition rate. a-Si:H films were fabricated by inductively coupled plasma CVD at 150 °C using 100% SiH₄ gas in pressure of 0.67 Pa and a flow rate of 50 sccm. The deposition rate was 0.8 nm/s.





Fig. 1. (Color online) Schematic apparatus for photo-induced carrier microwave absorption measurement system.

 1×10^{6} -Pa-H₂O vapor heat treatments at 180, 220, 260, and 300 °C for 1 and 3 h were applied to samples.

A 9.35-GHz-microwave transmittance measurement system with waveguide tubes was used to measure the photoinduced effective minority carrier lifetime, as shown in Fig. 1. The 9.35-GHz microwave was emitted by a fieldeffect-transistor (FET)-type oscillator. The microwave was introduced using a waveguide tube. There was a 1 mm narrow gap for the measurement of a sample wafer, which was able to be moved in the gap by an X-Y stage. Small holes were opened on walls of the waveguide tubes to place optical fibers for introducing light of 620-nm-light-emitting diodes (LEDs). The optical penetration depth of crystalline silicon was about $2.2\,\mu m$ at $620\,nm$,¹⁰ which was much lower than the substrate thickness. Carrier generation was limited in the surface region. Teflon plates were placed aslant in the waveguide tube to reflect and diffuse the incident light. Consequently, sample surfaces were uniformly illuminated at an intensity of 1.5 mW/cm² with LED light, which was pulsed for 2s, to precisely measure the effective minority carrier lifetime in the cases of light illumination at the top surface $\tau_{eff}(top)$ and at the rear surface $\tau_{\rm eff}$ (rear), as shown in Fig. 1. The microwaves transmitted through the samples were detected by high-speed diode rectifiers, and the signals were integrated in coincidence with the LED light 2-s pulse. The detection accuracy of the present system in the transmissivity was $\pm 0.1\%$. The integrated voltage was detected by a digital electrometer and analyzed to obtain $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ using carrier diffusion and annihilation theories.^{11,12}

3. Results and Discussion

The initial n-type silicon samples coated with 100-nm-thick

SiO₂ had high $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ about 1.2×10^{-3} s. $\tau_{eff}(top)$ was the same as $\tau_{eff}(rear)$. The silicon surfaces were well passivated by thermally grown SiO₂ layers. If the surface recombination velocities at the top surface S_{top} and rear surface S_{rear} are independent of light intensity, $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ are given according to carrier diffusion and annihilation theories as^{12,13}

$$\tau_{\rm eff}(\rm top) = \tau_{\rm b} \frac{\sqrt{\frac{D}{\tau_{\rm b}}} \left(1 - \exp\left(-\frac{d}{\sqrt{D\tau_{\rm b}}}\right)\right) \left(\sqrt{\frac{D}{\tau_{\rm b}}} + S_{\rm rear} + \left(\sqrt{\frac{D}{\tau_{\rm b}}} - S_{\rm rear}\right) \exp\left(-\frac{d}{\sqrt{D\tau_{\rm b}}}\right)\right)}{\left(\sqrt{\frac{D}{\tau_{\rm b}}} + S_{\rm rear}\right) \left(\sqrt{\frac{D}{\tau_{\rm b}}} + S_{\rm top}\right) - \left(\sqrt{\frac{D}{\tau_{\rm b}}} - S_{\rm top}\right) \left(\sqrt{\frac{D}{\tau_{\rm b}}} - S_{\rm rear}\right) \exp\left(-\frac{2d}{\sqrt{D\tau_{\rm b}}}\right)},$$

$$\tau_{\rm eff}(\rm rear) = \tau_{\rm b} \frac{\sqrt{\frac{D}{\tau_{\rm b}}} \left(1 - \exp\left(-\frac{d}{\sqrt{D\tau_{\rm b}}}\right)\right) \left(\sqrt{\frac{D}{\tau_{\rm b}}} + S_{\rm top} + \left(\sqrt{\frac{D}{\tau_{\rm b}}} - S_{\rm top}\right) \exp\left(-\frac{d}{\sqrt{D\tau_{\rm b}}}\right)\right)}{\left(\sqrt{\frac{D}{\tau_{\rm b}}} + S_{\rm rear}\right) - \left(\sqrt{\frac{D}{\tau_{\rm b}}} - S_{\rm rear}\right) \left(\sqrt{\frac{D}{\tau_{\rm b}}} - S_{\rm top}\right) \exp\left(-\frac{2d}{\sqrt{D\tau_{\rm b}}}\right)},$$

$$(2)$$

where *D* is the minority carrier diffusion coefficient, *d* is the substrate thickness, and $\tau_{\rm b}$ is the minority carrier bulk lifetime. A high effective lifetime means a low $S_{\rm top}$ and $S_{\rm rear}$. If $\tau_{\rm b}$ and $\tau_{\rm eff}$ are long enough, eqs. (1) and (2) can be simply approximated as

$$\tau_{\rm eff}({
m top}) \sim \tau_{\rm eff}({
m rear}) \sim {d \over S_{
m top} + S_{
m rear}} \,.$$
 (3)

Equations (1)–(3) indicate that high $\tau_{\rm eff}(\rm top)$ and $\tau_{\rm eff}(\rm rear)$ have almost the same value when $S_{\rm top}$ and $S_{\rm rear}$ are independent of light intensity. We previously revealed that coincidence between $\tau_{\rm eff}(\rm top)$ and $\tau_{\rm eff}(\rm rear)$ was obtained when the minority carrier diffusion length ($D^*\tau_{\rm eff}$)^{0.5} was longer than the thickness of the substrate.¹² In the present condition of hole minority carriers, $\tau_{\rm eff}(\rm top)$ and $\tau_{\rm eff}(\rm rear)$ have the same value when they are longer than 4×10^{-4} s. $S_{\rm top}$ and $S_{\rm rear}$ were therefore estimated to be about 22 cm/s for the initial n-type silicon samples.

When SiO₂ layers at the top surface of silicon substrates were removed, $\tau_{\rm eff}(top)$ was markedly decreased to about 1.6×10^{-5} s from the initial value of 1.5×10^{-3} s because the bare surface had a high recombination velocity. $\tau_{\rm eff}(rear)$ was also decreased to 1.2×10^{-4} s. Photo-induced carriers generated at the rear surfaces diffused toward the top surfaces and annihilated there. $S_{\rm top}$ was estimated from $\tau_{\rm eff}(top)$ and $\tau_{\rm eff}(rear)$ to be about 3000 cm/s.

Figure 2 shows $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ as a function of thickness of a-Si films formed by PECVD at 150 °C at the top surface. $\tau_{eff}(top)$ increased from 2.5×10^{-5} to 7.4×10^{-5} s as the film thickness increased from 3 to 50 nm. This means that a-Si:H played a role in surface passivation, and that thick a-Si:H was effective for surface passivation. We interpret that the increase in $\tau_{eff}(top)$ results from the increase in the ratio of surface coverage with increasing film thickness because it was not easy for very thin a-Si:H films to completely cover the crystalline silicon surface owing to mechanical stress.^{14,15)} An island-like a-Si:H domain would be formed on the surface under the conditions of 3- and 10-nm-thick a-Si:H film deposition. There would be a small



Fig. 2. $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ as a function of thickness of a-Si films formed at the top surface. Arrows present $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ for samples with as-SiO₂ removed at the top surface.

residual bare surface region, which decreases τ_{eff} . Our deposition conditions should be further improved to achieve complete surface coverage with thin a-Si:H films. On the other hand, τ_{eff} (rear) was hardly changed by a-Si:H film formation. The result of the independence of $\tau_{eff}(rear)$ with the thickness of a-Si:H films is strange. An increase in $\tau_{\rm eff}(top)$ should be associated with an increase in $\tau_{\rm eff}(rear)$ because photo-induced minority carriers diffuse from the top surface to the rear surface when the diffusion length of photo-induced minority carriers is longer than the substrate thickness of 520 µm with $\tau_{\rm b}$ above 2.3 × 10⁻⁴ s. Since the initial $\tau_{\rm eff}(top)$ was 1.5×10^{-3} s, much higher than $2.3 \times$ 10⁻⁴ s, photo-induced minority carriers were capable of diffusing from the top surface to the rear surface. We interpret that plasma damage occurred at both surfaces. The top surface would have been passivated by a-Si:H film formation, but simultaneously suffered from plasma damage. The rear surface would have only suffered from plasma damage and $\tau_{eff}(rear)$ would be decreased.

Figure 3 shows $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ as a function of heating temperature of 1×10^6 -Pa H₂O vapor heat treatment for 1 h for samples with a-Si:H film of 3 (a), 10 (b), and 50



Fig. 3. $\tau_{\rm eff}(top)$ and $\tau_{\rm eff}(rear)$ as a function of heating temperature of 1×10^6 -Pa H₂O vapor heat treatment for 1 h for samples with a-Si:H films of 3 (a), 10 (b), and 50 (c) nm thicknesses formed at the top surfaces.

(c) nm thickness formed at the top surfaces. $\tau_{eff}(top)$ increased to 8×10^{-5} – 1.6×10^{-4} s for samples with 3-nmthick a-Si:H films. 1×10^{6} -Pa H₂O vapor heat treatment improved the surface passivation at the top surface. $\tau_{eff}(rear)$ also increased to 2×10^{-4} - 3×10^{-4} s. H₂O vapor heat treatment at high temperatures of 260 and 300 °C resulted in high $\tau_{eff}(top)$ and $\tau_{eff}(rear)$, as shown in Fig. 3(a). Plasma damage was probably reduced for both surfaces. On the other hand, $\tau_{\rm eff}(top)$ increased from 1.1×10^{-4} to $3.0 \times$ 10^{-4} s as the temperature of 1×10^{6} -Pa H₂O vapor heat treatment decreased from 300 to 180 °C for samples with 10-nm-thick a-Si:H films. However, $\tau_{eff}(rear)$ was almost leveled off between 2.0×10^{-4} and 2.2×10^{-4} s. This indicates that effective passivation occurred in the case of light illumination at the top surface for low temperature conditions at 180 and 220 °C of 1.0×10^6 -Pa H₂O vapor heat treatment, as shown in Fig. 3(b). A high $\tau_{eff}(top)$ of 1.15×10^{-3} s was observed for samples with 50-nm-thick a-Si films treated with 1×10^6 -Pa H₂O vapor heat treatment at 180 °C. $\tau_{\rm eff}(top)$ markedly increased from 5.4 × 10⁻⁴ to 1.15×10^{-3} s as the temperature of H₂O vapor heat treatment decreased from 300 to $180 \,^{\circ}\text{C}$. $\tau_{eff}(\text{rear})$ also increased from 4.0×10^{-4} to 9.3×10^{-4} s as the temperature of 1×10^6 -Pa H₂O vapor heat treatment decreased from 300 to 180 °C.

It is interesting that very effective passivation occurred in the case of light illumination at the top a-Si:H surface for low temperature conditions at 180 and 220 °C of 1.0×10^6 -Pa H₂O vapor heat treatment, as shown in Fig. 3(c) for samples coated with 50-nm-thick a-Si films. The experimental results of Fig. 3 revealed that samples coated with 50-nm-thick a-Si:H at the top surface had a $\tau_{\rm eff}$ (top) value



Fig. 4. (Color online) $\tau_{\rm eff}$ as a function of light intensity in the cases of four different light illumination modes shown in insets for samples with 50-nm-thick a-Si:H annealed at 180 °C with 1×10^{6} -Pa H₂O vapor heat treatment for 1 h. The light intensity of bias light was 0.25 mW/cm².

about 30% higher than the $\tau_{\rm eff}$ (rear) value. Although those values were longer than 4×10^{-4} s, in the cases of 1.0×10^{6} -Pa H₂O vapor heat treatment at 180 and 220 °C. This result indicates a different physics of passivation from the carrier annihilation story with independent recombination velocities of light intensity as discussed above.

To clarify this phenomenon experimentally, we measured $\tau_{\rm eff}$ for the following four different types of light illumination for samples with 50-nm-thick a-Si:H annealed at 180 °C, as shown in Fig. 4. τ_{eff} was measured by light illumination at the top a-Si:H films at an intensity ranging from 0 to $1.5 \,\mathrm{mW/cm^2}$ with the rear surface in the dark, case I (open circles and solid line) and with a bias light at $0.25 \,\mathrm{mW/cm^2}$ illuminated at the rear surface, case II (open squares and solid line). τ_{eff} was also measured by light illumination at the rear SiO2 films at an intensity ranging from 0 to $1.5 \,\mathrm{mW/cm^2}$ with the top surface in the dark, case III (solid circles and dashed line) and with a bias light at $0.25 \,\mathrm{mW/cm^2}$ illuminated at the a-Si:H films, case IV (solid squares and dashed line). A high τ_{eff} ranging from 1.1×10^{-3} to $1.15\times10^{-3}\,\mathrm{s}$ was observed in the cases of light illumination at the a-Si:H films of cases I, II, and IV. On the other hand, low $\tau_{\rm eff}$ ranging from 8.0×10^{-4} to 8.5×10^{-4} s were observed in the cases III and II for the top a-Si:H surface in the dark. Those results clearly show that light-illuminated a-Si:H had a role in the improvement of surface passivation and increase in $\tau_{\rm eff}$.

Changes in photo-induced carrier density per unit area were measured as a function of time in the case of 50-nmthick a-Si:H samples with 1.0×10^{6} -Pa H₂O vapor heat treatment at 180 °C for 1 h, as shown in Fig. 5. The samples were illuminated at the top a-Si:H surface at 1.5 mW/cm^{2} for an initial 10 s and continuously illuminated at the rear surface [curve (a)], illuminated at the rear surface at 1.5 mW/cm^{2} for initial 10 s and continuously illuminated at the top a-Si:H surface [curve (b)], only continuously illuminated at the top a-Si:H surface at 1.5 mW/cm^{2} with



Fig. 5. Changes in photo-induced carrier density per unit area for four different light illumination modes for samples with 50-nm-thick a-Si:H annealed at 180 °C with 1×10^6 -Pa H₂O vapor heat treatment for 1 h. The light intensity was 1.5 mW/cm^2 .

keeping the rear surface dark [curve (c)], and only continuously illuminated at the rear surface at 1.5 mW/ cm^2 with keeping the top surface dark [curve (d)]. The carrier density for curve (a) was the same as that for curve (b) until 10 s. It was also the same as the sum of the carrier densities for curves (c) and (d). It rapidly decreased to the level of curve (c) immediately after light termination at 10s. It subsequently decreased gradually to that for curve (d). This clearly shows that light-illuminated a-Si:H films maintained good passivation for a long time with a time constant of about 3 s. On the other hand, in the case of curve (b), the carrier density rapidly decreased to that for curve (c) at 10s and subsequently leveled off. These experimental results of Figs. 4 and 5 clearly indicate that the illumination of a-Si:H has an unique role in improving surface passivation. Although we have unfortunately not established a physical theory to explain this interesting phenomenon yet, we give a possible model on the basis of our experiences.

The measurement of optical absorption revealed that 50nm-thick a-Si:H films absorbed 3% of the incident 620 nm light. Hole and electron carriers were generated in the a-Si:H films. We believe that hole carriers were trapped in the a-Si:H films, which were annealed with high-pressure H₂O vapor heat treatment. Hole trapping bent the silicon band at the surface region according to the strict charge neutrality rule given by the Euler condition and caused an electron accumulation region at the surface resulting in the decrease in S_{top} and in the increase in $\tau_{eff}(top)$. We already confirmed that the density of surface trapped states and plasma damage were decreased by high-pressure H₂O vapor heat treatment.^{6,7)} Moreover, oxygen atoms were introduced into the a-Si:H films during high-pressure H₂O vapor heat treatment.¹⁶⁾ Si-O bondings were substantially formed by breaking Si–H bonding. We also reported that unstable SiO_x played a role in hole trapping states.^{17,18} We interpret that our condition of high-pressure H₂O vapor heat treatment for 1h formed unstable Si-O bondings trapping hole carriers generated by light illumination, although interface traps and plasma damage must have been reduced. Electron carriers generated by light illumination would be diffused out into crystalline silicon because the a-Si/c-Si structure was conductive. A $\tau_{eff}(top)$ of $1.15 \times 10^{-3} \, s$ and a $\tau_{eff}(rear)$ of 9.3×10^{-4} s were obtained for high-pressure H₂O vapor heat treatment at 180 °C. Those $\tau_{\rm eff}$ values resulted the $S_{\rm top}$ decreasing from 28 (dark) to 18 cm/s when the a-Si:H film was illuminated by 620-nm light. However, the band structure of the a-Si:H/Si layered structure in a dark field was not clear for samples annealed with high-pressure H₂O vapor for 1 h. a-Si:H and crystalline silicon have very different band gaps. The band offset between the a-Si:H/Si interface has also been precisely investigated.¹⁹⁻²¹⁾ It has been reported that the difference in their conduction band edges can be small, while the difference in their valence band edges can be large. However, our present samples had no metal electrodes and no metal oxide semiconductor structure. τ_{eff} was measured in a floating condition with no standard potential voltage. We therefore did not have information of the initial band bending at the silicon surface. When the band bent upward associated with the electrondepletive surface region caused by strong dipole moments induced by the band offset in the dark field, the band would be bent downward associated with electron accumulated region by a positive-charge-induced field effect, when light was illuminated at the top s-Si:H surface. S_{top} therefore decreased according to standard carrier annihilation theory.¹³⁾ If there was still a substantial density of interface traps at the mid gap, the band would be also bent upward associated with the electron-depletive surface region in the dark field. The band would be also bent downward in this case when light was illuminated at the top s-Si:H surface.

From our experiences, high-pressure H₂O vapor heat treatment for a long time and at high temperatures effectively decreased the density of dangling bonds in silicon and at its surface, and the treatment changes unstable SiO_x (x < 2) to stable SiO_2 .^{17,18}) We applied 1.0×10^6 -Pa H₂O vapor heat treatment for a long time of 3 h. Figure 6 shows $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ as a function of heating temperature of 1×10^6 -Pa H₂O vapor heat treatment for 3 h for samples with a-Si:H films of 3 (a), 10 (b), and 50 (c) nm thicknesses formed at the top surfaces. No substantial improvement in τ_{eff} was observed for samples with 3-nmthick a-Si:H films, as shown in Fig. 6(a) compared with the case of H₂O vapor heat treatment for 1 h, as shown in Fig. 3(a). Slight increases in τ_{eff} were observed for samples with 10-nm-thick a-Si:H films for 1×10^{6} -Pa H₂O vapor heat treatment at 180 and 300 °C. However, $\tau_{eff}(top)$ was still larger than $\tau_{\rm eff}$ (rear) in the cases of high-pressure H₂O vapor heat treatment at 180 and 300°C, as shown in Fig. 6(b). This indicates that the light-induced passivation was still effective in the case of 3h H₂O vapor heat treatment. On the other hand, $\tau_{eff}(top)$ and $\tau_{eff}(rear)$ were much increased by 1×10^{6} -Pa H₂O vapor heat treatment for samples with 50-nm-thick a-Si:H films. In particular, $\tau_{\rm eff}(top)$ was increased to 1.2×10^{-3} s and $\tau_{\rm eff}(tear)$ was increased to 1.4×10^{-3} by 1×10^{6} -Pa H₂O vapor heat treatment at 300 °C for 3 h, as shown in Fig. 6(c). Those τ_{eff} were close to the initial τ_{eff} of 1.5×10^{-3} s for the initial samples with both surfaces coated with 100-nm-thick thermally grown SiO₂. Both silicon surfaces were well passivated with a combination of a-Si:H film formation with high-pressure H₂O vapor heat treatment at 300 °C.

In the case of 50-nm-thick a-Si:H samples with 1.0×10^6 -Pa H₂O vapor heat treatment at 300 °C for 3 h,



Fig. 6. $\tau_{\rm eff}({\rm top})$ and $\tau_{\rm eff}({\rm rear})$ as a function of heating temperature of 1×10^6 -Pa H₂O vapor heat treatment for 3 h for samples of a-Si:H films of 3 (a), 10 (b), and 50 (c) nm thicknesses formed at the top surfaces.

H₂O vapor heat treatment [°C]



Fig. 7. Changes in photo-induced carrier density per unit area for four different light illumination modes for samples with 50-nm thick a-Si:H annealed at 300 °C with 1×10^6 -Pa H₂O vapor heat treatment for 3 h. The light intensity was 1.5 mW/cm^2 .

Fig. 7 shows the changes in photo-induced carrier density per unit area measured with time for light illumination to the a-Si:H films at 1.5 mW/cm^2 for an initial 10 s and continuous illumination to the rear surface [curve (a)], light illumination to the rear surface at 1.5 mW/cm^2 for initial 10 s and continuous illumination to the top a-Si:H surface [curve (b)], continuous light illumination only to the top a-Si:H surface at 1.5 mW/cm^2 [curve (c)], and continuous light illumination only to the rear surface at 1.5 mW/cm^2 [curve (d)]. The carrier density for curve (a) was the same as that for curve (b) until 10 s. It was also the same as the sum of the carrier densities for curves (c) and (d). It rapidly decreased immediately after light termination at 10s and leveled off at the same value of the photo-induced carrier density per unit area of curve (d). No light-induced passivation was observed in the present time range for samples coated with 50-nm-thick a-Si:H in the case of highpressure H₂O vapor heat treatment for 3 h. We believe that a-Si:H films were oxidized well by high-pressure H₂O vapor heat treatment for 3 h, and that stable SiO₂ with no capablity of trapping holes was formed in a-Si:H. Our previous study revealed that oxygen incorporation by high-pressure H₂O vapor heat treatment for 3h was small 1 at. % on average at most, and that the optical band gap was hardly changed by the treatment.¹⁶⁾ These findings suggest that SiO_2 was formed as small cluster structure in a-Si:H films. Our model is therefore that high-pressure H₂O vapor heat treatment for 3h well passivated the a-Si:H/c-Si interfaces and formed stable SiO₂ clusters, which did not trap photo-induced hole carriers, so that no light-induced passivation was observed. Light-induced passivation probably occurred associated with unstable SiO_x bonds trapping hole carriers. Although our model has not been experimentally established, the experimental results of Figs. 3-5 clearly demonstrate interesting light induced passivation in the case of illumination of a-Si:H. Those results stimulate further material investigation of stable species to be incorporated to cause effective light-induced passivation by their photo-induced holes or electrons.

4. Conclusions

We investigated the surface passivation of n-type crystalline silicon coated with a-Si:H films at the top surface, which were annealed in 1.0×10^6 Pa H₂O vapor for 1 and 3 h. The 9.35-GHz microwave transmittance measurement system with 620-nm light illumination at the top a-Si:H films as well as the rear SiO₂ films was used to measure τ_{eff} . High τ_{eff} values of 1.6×10^{-4} , 3.0×10^{-4} , and 1.15×10^{-3} s at maximum were achieved by 1.0×10^6 Pa H₂O vapor heat treatment between 180 and 300 $^\circ C$ for 1 h for samples with the top surfaces formed with a-Si:H films with thicknesses of 3, 10, and 50 nm, respectively. Light-induced enhancement in τ_{eff} was observed when light was illuminated to the 50nm-thick a-Si:H films. τ_{eff} increased by about 30% when the a-Si:H films were illuminated. The enhancement in τ_{eff} was maintained for a long time with a time constant of 3 s, when the light was off. Electrical field effect passivation was probably caused by hole trapping at SiO_x clusters formed in the a-Si:H films by 1.0×10^6 Pa H₂O vapor heat treatment for 1 h. On the other hand, heat treatment at 300 °C with 1.0×10^6 Pa H₂O vapor for 3 h increased $\tau_{eff}(top)$ to 1.2×10^{-3} and $\tau_{\rm eff}$ (rear) to 1.4×10^{-3} s, which were close to the initial τ_{eff} of 1.5×10^{-3} s for samples with both surfaces coated with 100-nm-thick thermally grown SiO₂. No enhancement in τ_{eff} was observed when a-Si:H films were illuminated in that case.

Acknowledgements

This work was partly supported by Grants-in-Aid for Science Research C (Nos. 22560292 and 23560360) from the Ministry of Education, Culture, Sports, Science and Technology of Japan, and Mikiya Industrial and Economic Research Foundation.

- A. Froitzheim, K. Brendel, L. Elstner, W. Fuhs, K. Kliefoth, and M. Schmidt: J. Non-Cryst. Solids 299–302 (2002) 663.
- S. Dauwe, J. Schmidt, and R. Hezel: Proc. 29th IEEE Photovoltaic Specialists Conf., 2002, p. 1246.
- 3) S. D. Wolfa and M. Kondo: Appl. Phys. Lett. 90 (2007) 042111.
- M. Taguchi, A. Terakawa, E. Maruyama, and M. Tanaka: Prog. Photovoltaics 13 (2005) 481.
- M. Taguchi, E. Maruyama, and M. Tanaka: Jpn. J. Appl. Phys. 47 (2008) 814.
- M. Hasumi, J. Takenezawa, T. Nagao, and T. Sameshima: Jpn. J. Appl. Phys. 50 (2011) 03CA03.
- 7) T. Sameshima and M. Satoh: Jpn. J. Appl. Phys. 36 (1997) L687.
- T. Sameshima, M. Satoh, K. Sakamoto, K. Ozaki, and K. Saitoh: Jpn. J. Appl. Phys. 37 (1998) 4254.
- E. Takahashi, Y. Nishigami, A. Tomyo, M. Fujiwara, H. Kaki, K. Kubota, T. Hayashi, K. Ogata, A. Ebe, and Y. Setsuhara: Jpn. J. Appl. Phys. 46 (2007) 1280.
- E. D. Palk: Handbook of Optical Constants of Solids (Academic Press, London, 1985) p. 547.

- 11) T. Sameshima, H. Hayasaka, and T. Haba: Jpn. J. Appl. Phys. 48 (2009) 021204.
- T. Sameshima, T. Nagao, S. Yoshidomi, K. Kogure, and M. Hasumi: Jpn. J. Appl. Phys. 50 (2011) 03CA02.
- 13) A. S. Grove: Physics and Technology of Semiconductor Devices (Wiley, New York, 1967) Chap. 5.
- 14) H. Fujiwara, Y. Toyoshima, M. Kondo, and A. Matsuda: Phys. Rev. B 60 (1999) 13598.
- 15) H. Fujiwara and M. Kondo: J. Appl. Phys. 101 (2007) 054516.
- 16) T. Sameshima, H. Hayasaka, M. Maki, A. Masuda, T. Matsui, and M. Kondo: Jpn. J. Appl. Phys. 46 (2007) 1286.
- 17) K. Sakamoto and T. Samaehim: Jpn. J. Appl. Phys. 39 (2000) 2492.
- 18) H. Watakabe, T. Sameshima, T. Strutz, T. Oitome, and A. Kohno: Jpn. J. Appl. Phys. 44 (2005) 8367.
- 19) J. Tersoff: Phys. Rev. B 30 (1984) 4874.
- 20) C. Tejedor and F. Flores: J. Phys. C 11 (1978) L19.
- 21) T. F. Schulze, L. Korte, F. Ruske, and B. Rech: Phys. Rev. B 83 (2011) 165314.