The influence of the hot wire temperature on the crystallization of $\mu\text{c-Si:H}$ films prepared by hot wire-assisted-ECR-CVD

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ABSTRACT

We have constructed a hot wire assisted ECR CVD system to prepare a-Si:H and $\mu\text{c-Si:H}$ films. The effect of hot wire temperature on crystallization of a-Si:H films is studied in the films prepared by these system. The crystalline fraction and crystalline grain size are analyzed by Raman scattering. The hydrogen content and the bonding scheme of hydrogen were measured by Fourier transform infrared (FTIR) spectroscopy. At low hot wire temperature, about 20 at.% hydrogen is included in the film. With increasing the hot wire temperature, the total content of the hydrogen, SiH_2 and SiH decrease and the microcrystalline phase appears. It is found from the ratio of the crystalline TO peak to the total area of the TO peak of the Raman scattering spectra that the volume fraction of the crystalline phase increases with increasing the hot wire temperature. The crystalline peak has a tendency to shift toward the higher wavenumber with increasing the hot wire temperature, suggesting that the grain size increases with increasing the hot wire temperature.

INTRODUCTION

In recent years the applications of hydrogenated microcrystalline silicon ($\mu\text{c-Si:H}$) films, grown at low substrate temperature, in thin film transistors (TFT) for use in active matrix liquid crystal displays, p-i-n solar cells and light emitting devices (LED) have been intensively investigated. Light-induced degradation of a-Si:H based solar cells has been a major impediment in the development of thin film amorphous silicon technology. The stability of a-Si:H films is related to the amorphous network structure, whereas the improvement of amorphous network may finally lead to the formation of micro or nanocrystalline silicon ($\mu\text{c-}$ or nc-Si), which shows no noticeable light-induced degradation, but a poor photosensitivity. It has been confirmed in recent years that hydrogenated silicon films deposited in the regime just above the phase transition from amorphous to crystalline state could gain both the fine photoelectronic properties like a-Si:H and high stability like $\mu\text{c-Si:H}$ [1-6]. The demand of these applications is not only a high quality but also a high deposition rate on the preparation. Although the plasma-enhanced CVD is commonly used for the preparation of $\mu\text{c-Si:H}$, the electron cyclotron resonance (ECR) CVD and hot wire (HW) CVD have also been studied as promising methods to obtain a high rate deposition at low substrate temperature. We have constructed a hot wire assisted ECR CVD system to prepare a-Si:H and $\mu\text{c-Si:H}$ films.

In this paper, a-Si:H and $\mu\text{c-Si:H}$ films have been deposited at various hot wire temperature in a hot wire assisted MWECR CVD apparatus. The effect of hot wire temperature on crystallization of a-Si:H films is studied in the films prepared by the above system.

Fig.1. Schematic diagram of the experimental apparatus.

EXPERIMENTAL DETAILS

A-Si:H and $\mu\text{c-Si:H}$ films were prepared using hot wire assisted MWECD CVD system. The schematic diagram of the experimental apparatus is shown in Figure 1. The frequency of the microwave was 2.45GHz, the power was 555 W and the intensity of magnetic field in resonance section was 0.0875T. In order to increase the activity of the particles, we used hot wire assisted technique. The tungsten filaments of 0.5 mm thick and 350 mm long, located at about 60 mm from the substrate. The voltage in the filaments was controlled by a booster; the temperature of filament was measured by an infra-red pyrometer. The substrate temperature was 473K, pressure was 2 Pa. The flow flux is 20sccm, and the hydrogen dilution ratio is 1:4. The films were deposited on single crystal Si wafers and Corning-7059 glasses respectively. The film thickness was measured by a TENCOR alpha-step profilometer. The thickness of films was about 1 μm . The crystalline volume fraction and crystalline grain size are analyzed by Raman scattering. The hydrogen content and the bonding scheme of hydrogen were measured by Fourier transform infrared (FTIR) spectroscopy.

RESULTS AND DISCUSSION

Fig. 2 Variation of deposition rate as a function of hot wire temperature of the films.

Figure 2 shows the variation of deposition rate as a function of the hot wire temperature. As seen in the figure, the deposition rate increases from 19.8 to 26.7 Å/s, as the HW temperature increases up to 1600°C. The deposition rates in the present work are much higher than those by the ECR CVD [7] and those by the HW CVD in the low HW temperature range [8].

Fig. 3 (a) FTIR absorption

Fig. 3 (b) Hydrogen content of the

The FTIR absorption spectra of the films are shown in Figure 3(a). Two peaks of the stretching mode of SiH and SiH₂ were observed at 2000cm⁻¹ and 2100cm⁻¹, respectively. Two Gaussian peaks were used to fit the observed spectra. The amount of hydrogen bonded in SiH and SiH₂ configurations (C_{SiH} and C_{SiH2}) was calculated from the area of each assigned peak and the total content of hydrogen (C_H) was the sum of C_{SiH} and C_{SiH2}. The hydrogen contents are shown in Figure 3(b). With increasing the hot-wire temperature, the total hydrogen content and the SiH₂ content decrease, but the SiH content first increases and then decreases, and above the 1450°C, the SiH content almost do not change.

Transverse optic (TO) modes of Raman spectra obtained from the μ c-Si:H films prepared at various hot wire temperature are shown in Figure 4. As the hot-wire temperature increased, the c-Si TO peak (at around 520cm⁻¹) appears and increases. From Fig.4, it can be found that the crystalline peak has a tendency to shift toward the higher wavenumber with increasing the hot wire temperature, suggesting that the grain size increases with increasing the hot wire temperature.

Fig. 4 Raman spectra of the films deposited with different hot-wire

The grain size and the volume crystalline fraction of the films can be deduced by using the TO mode. Assuming a confinement model, the Raman shift is related with the size L of the crystallite as the formula (1) [9]:

$$\Delta\omega = \omega(L) - \omega_0 = -A(a/L)^\gamma \quad (1)$$

where $\omega(L)$ is the frequency of the Raman phonon with size L , ω_0 is the frequency of the optical phonon at the zone center, and a is the lattice constant of Si. The parameters A and γ used to describe the vibrational confinement are 47.41cm⁻¹ and 1.44, respectively. From Fig.4, $\Delta\omega$ is found to be 4.5cm⁻¹, 2.5cm⁻¹ and 1.5cm⁻¹ for samples that with the hot wire temperature of 1450°C, 1500°C and 1600°C, respectively. Then we obtain the corresponding grain size of 2.8nm,

4.2nm and 6.0nm, respectively. The increase in the grain size may be affected by the change in the structural disorder in the amorphous phase which is correlated to the SiH bonding characteristics of the film.

In order to obtain the crystalline volume fraction X_c , the Raman spectra of the TO mode were deconvolved into three components [10]. The lower wavenumber component (a) at around 480 cm^{-1} due to a-Si, the higher wavenumber component (c) at around 520 cm^{-1} due to c-Si and the intermediate component (b) at around 510 cm^{-1} due to the bond dilation at the grain boundaries. The crystalline volume fraction was deduced from the formula (2):

$$X_c = (I_b + I_c) / (I_a + I_b + I_c) \quad (2)$$

where I_a , I_b and I_c denote the intensity of the integrated intensity of the three components, respectively. It is found that the X_c of the $\mu\text{c-Si:H}$ films is 53.6%, 60.3% and 78.2%, which hot-wire temperature of 1450°C , 1500°C and 1600°C , respectively.

The results mentioned above are quite different from those of the films synthesized by either HW CVD or ECR CVD method. The coverage over the growing surface with atomic hydrogen is required for the forming of micro crystallites because the adsorbed SiH_3 can migrate to find a low energy site for crystallization when the growing surface is covered with hydrogen atoms [11]. In the case of HW CVD, the atomic hydrogen is supplied by the decomposition of SiH_4 on the hot wire. Sufficient amounts of hydrogen atoms for the surface coverage as well as an amount of adsorbed SiH_3 to keep the film deposition are attained when the HW temperature is high, e.g. 2100°C [8]. In the case of ECR CVD, however, a very high dilution rate of H_2/SiH_4 is used for preparation of the $\mu\text{c-Si:H}$. Moreover, a high microwave power P_{mw} is needed to get the surface coverage with hydrogen atoms and a high substrate temperature T_s is needed to make adsorbed SiH_3 migrate on the growing surface, for example, $P_{\text{mw}} \geq 540\text{W}$ and $T_s \geq 250^\circ\text{C}$ [7]. We can prepare $\mu\text{c-Si:H}$ at lower HW temperatures than by the HW CVD and at lower dilution ratio, P_{mw} and T_s than by the ECR CVD.

It is reported for a-Si:H films prepared by HW CVD that the deposition rate dramatically increases and the hydrogen content increases with the HW temperature [8]. In the present work, however, the deposition rate increases slightly with the HW temperature as shown in Fig. 2 and the hydrogen content decreases with the HW temperature as shown in Fig. 3(b). The rather high deposition rate in the a-Si:H film deposited with $T_w \leq 1400^\circ\text{C}$ in present work compared to ref. 8 can be attributed to the enhancement by the effect of ECR. The atomic hydrogen created in the ECR chamber may enhance the decomposition of SiH_4 to increase the decomposition rate. Both the films prepared without operating HW and those prepared with $T_w = 1400^\circ\text{C}$ are in the amorphous phase. But the latter films have a higher concentration of hydrogen atoms in the monohydride phase in spite of the lower concentration of total bonded hydrogen atoms than the former films as shown in Fig. 3(b). This suggests that not only the hydrogen atoms originating from the ECR plasma but also those created by HW contribute to yield a good quality a-Si:H. At higher HW temperature ($\geq 1450^\circ\text{C}$), micro crystallites appear in the films. The excess atomic hydrogen may attack the growing surface of the film and have an etching effect and so the deposition rate decreases, but the overall deposition rate increases under the competition between the decomposition and the etching. Because of the etching effect, the hydrogen content in the films tends to decrease with increasing the HW temperature as shown in Fig. 3(b).

CONCLUSIONS

The effect of hot wire temperature on the properties of $\mu\text{c-Si:H}$ was studied. The deposition rate of the samples dependent on the hot wire temperature. The total hydrogen content of the films showed a tendency to decrease with the hot wire temperature increasing. The hydrogen content of SiH first increases and then decreases with the hot wire temperature increasing. And from 1450°C , microcrystalline silicon phase appears. With increasing the hot wire temperature, crystalline volume fraction and crystalline grain size increase. Microcrystallization to the Si:H network has been easily obtained from the hot wire assisted ECR CVD method at a relatively low power, low substrate temperature and low hot wire temperature.

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