

Deposition of highly stable cluster-free a-Si:H films using a fast gas flow multi-hollow discharge plasma CVD method

高速ガス流マルチホロープラズマCVD法を用いた
高光安定クラスタフリーa-Si:H膜の作製

Kosuke Hatozaki¹, Kenta Nakahara¹, Yuuji Hashimoto¹, Takeaki Matsunaga¹,
Daisuke Yamashita¹, Hidefumi Matsuzaki¹, Hyunwoong Seo¹, Kunihiro Kamataki²,
Giichiro Uchida¹, Naho Itagaki¹, Kazunori Koga¹ and Masaharu Shiratani¹
波戸崎浩介, 中原賢太¹, 橋本優史¹, 松永剛明¹, 山下大輔¹, 松崎秀文¹,
徐鉉雄¹, 鎌滝晋礼², 内田儀一郎¹, 板垣奈穂^{1,3}, 古閑一憲¹, 白谷正治¹

¹Graduate School of Information Science and Electrical Engineering, Kyushu University

²Center for Research and Advancement in Higher Education Kyushu University

^{1,2}744 Motooka, Nishi-ku, Fukuoka 819-0395, Japan

³PRESTO, Japan Science and Technology Agency, 5 Sanbancho, Chiyoda-ku, Tokyo 102-0075, Japan

^{1,2}九州大学大学院システム情報科学府 〒819-0395 福岡市西区元岡744

³JSTさきがけ 〒102-0075 東京都千代田区三番町5

We have deposited cluster-free a-Si:H films using a fast gas flow multi-hollow discharge plasma CVD method. To evaluate their quality as an I layer of PIN solar cells, we have measured Fill Factor (FF) of N-type c-Si/a-Si:H/Ni Schottky cells of such cluster-free a-Si:H films. Our films deposited at a rate of 0.51 nm/s show high initial FF of 0.556, high stable FF of 0.523, and quite low light induced degradation ratio of 5.93 %.

1. Introduction

Light-induced degradation of hydrogenated amorphous silicon (a-Si:H) has been an important issue for a-Si:H solar cells, because light exposure initially causes a reduction of the conversion efficiency of the cells due to the degradation [1]. During the deposition, amorphous silicon particles (clusters) in a size range below 10 nm are generated in the plasmas and they deposit into the films [2]. For conventional plasma CVD of a-Si:H films, clusters are generated in discharges and deposit into the films, and degrade efficiency and stability of the cells using such films. Reducing incorporation of the clusters is important for realizing highly stable a-Si:H solar cells of high efficiency [3]. It motivated us to deposit cluster-free a-Si:H films to realize stable a-Si:H solar cells of a high efficiency.

Recently, we have deposited highly stable a-Si:H films of $4.7 \times 10^{15} \text{ cm}^{-3}$ in stabilized defect density at a rate of 3 nm/s at substrate temperature $T_s = 250^\circ\text{C}$ by suppressing incorporation of clusters into the films [4,5]. The films show high stability against light exposure. In this study, we have deposited cluster-free a-Si:H films using the SiH_4 multi-hollow discharge plasma CVD method [4]. We report performance of Schottky solar cells using such

cluster-free a-Si:H films.

2. Experimental

Experiments were carried out using a multi-hollow discharge plasma CVD reactor as shown in Fig. 1. Multi-hollow powered and two grounded electrodes had holes of 5mm in diameter were placed in a stainless-steel vessel. SiH_4 gas was fed from a gas inlet at a flow rate of 150 sccm and was pumped out through the electrodes. The gas pressure was 0.5 Torr. The discharge frequency and power were 60 MHz and 40 W, respectively. Cluster-free a-Si:H films were deposited on a N-type c-Si substrate set in the upstream region

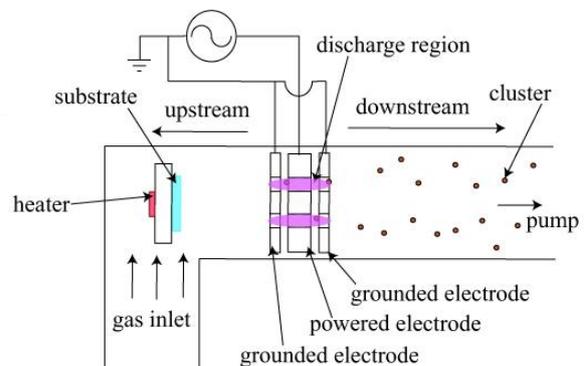


Fig.1. Multi-hollow discharge plasma CVD reactor.

from the discharges, because clusters generated in the discharges were driven toward the downstream region by gas flow. We deposited 605 nm films of a-Si:H at $T_s = 250^\circ\text{C}$. The deposition rate was 0.51 nm/s.

FF was evaluated irradiating N-type c-Si/a-Si:H/Ni Schottky cells by irradiation of an AM 1.5 solar spectrum light of 100 mW/cm^2 (1 SUN) at $T_s = 30^\circ\text{C}$. The stabilized FF was measured after 6 hours 40 min light soaking under conditions of 2.7 SUN irradiation and $T_s = 50^\circ\text{C}$.

3. Results and Discussion

Initial and stabilized J-V and P-V characteristic of the Schottky cell are shown in Fig. 2 and Table I. The fill factor (FF), maximum power (P_{max}), open circuit voltage (V_{oc}), short circuit current density (J_{sc}), series resistance (R_s), and parallel resistance (R_{sh}) are obtained from J-V curves of the Schottky cells. The initial and stabilized FF values

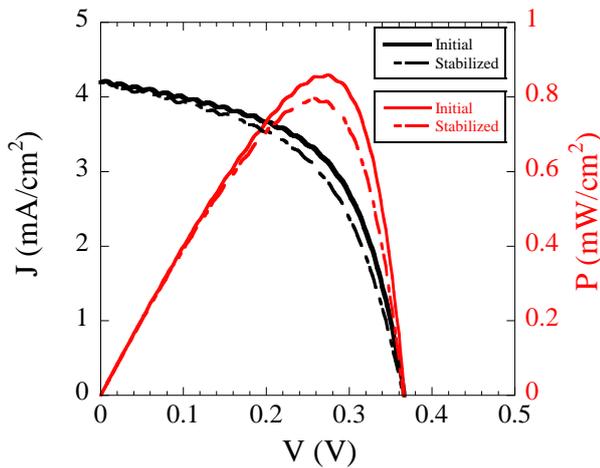


Fig. 2 Initial and stabilized J-V characteristic of Schottky solar cell. Conditions: SiH_4 150sccm, 0.5Torr, Frequency 60MHz, T_s 250°C , Electrode power 40W, 6h40min light soaking under conditions of 2.7 SUN and T_s 50°C .

Table I. Initial and stabilized characteristics of Schottky cell (FF, P_{max} , V_{oc} , J_{sc} , R_s , and R_{sh}) and degradation ratios of FF and P_{max} values

Characteristics of Schottky cell	Initial	Stabilized	Degradation ratio
FF	0.556	0.523	5.93%
P_{max} (mW/cm^2)	0.859	0.797	7.21%
V_{oc} (V)	0.37	0.37	
J_{sc} (mA/cm^2)	4.21	4.17	
R_s ($\Omega \text{ cm}^2$)	15.5	19.6	
R_{sh} ($\Omega \text{ cm}^2$)	524	466	

of Schottky cell are 0.556 and 0.523, respectively. This degradation ratio is 5.93%. Initial P_{max} of the cell was 0.859 mW/cm^2 . Stabilized P_{max} was 0.797 mW/cm^2 which was 7.21% lower than the initial one. Light soaking has little influence on V_{oc} and the initial and stabilized V_{oc} are 0.37 V. The initial and stabilized J_{sc} were 4.21 mA/cm^2 and 4.17 mA/cm^2 , respectively. These results suggest that a-Si:H films with little light induced degradation can be deposited by suppressing the cluster incorporation.

4. Conclusions

We have examined characteristics of N-type c-Si/a-Si:H/Ni Schottky cells using cluster-free a-Si:H films deposited by the multi-hollow discharge plasma CVD method. Our films deposited at a rate of 0.51 nm/s show high initial FF of 0.556, high stable FF of 0.523, and quite low light induced degradation ratio of 5.93%. These results indicate that our method is useful to fabricate highly stable a-Si:H solar cells.

Acknowledgements

This work was partly supported by the New Energy and Industrial Technology Development Organization (NEDO) and Photovoltaic Power Generation Technology Research Association (PVTEC).

References

- [1] D.L. Staebler and C.R. Wronski, *Appl. Phys. Lett.* **31** (1977) 292.
- [2] K. Koga, Y. Matsuoka, K. Tanaka, M. Shiratani, and Y. Watanabe, *Appl. Phys. Lett.* **77** (2000) 196.
- [3] K. Koga, N. Kaguchi, K. Bando, M. Shiratani, and Y. Watanabe, *Rev. Sci., Instrum.* **76** (2005) 113-501.
- [4] W. M. Nakamura, H. Matsuzaki, H. Sato, Y. Kawashima, K. Koga, M. Shiratani, *Suf. Coat. Technol.* **205** (2010) S241.
- [5] K. Koga, N. Kaguchi, K. Bando, and M. Shiratani, Y. Watanabe, *Rev. Sci., Instrum.* **76** (2005) 113501.