Dissolution Kinetics in Chemically Amplified EUV Resist

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Polymer structure effects on the dissolution kinetics and deprotection reaction were investigated to understand inherent extreme ultraviolet (EUV) resist characteristics. The effect of protecting groups and protecting ratio on activation energy were observed. Also, the protecting ratio effect on dissolution kinetics was observed. The speed of TMAH penetration into each resist film was different for each resist as the concept of the "penetration zone". This indicated that the dominant rate-determining step in resists is TMAH penetration into the resist film. Thus, the dissolution rates were affected by changing the protection ratio of polymer. Moreover, the preliminary evaluation of resist profile for EUV chemically amplified resists was performed by using PROLITH and parameters obtained with EUV exposure tool. Furthermore, the resist patterns were produced by electron beam and the SEM observation results were compared with the PROLITH simulation results. It is important for EUV lithography to select appropriate protecting group and protecting ratio.

Keywords: EUV lithography, chemically amplified resist, dissolution kinetics, deprotection reaction,

1. Introduction

Beyond the 16 nm node, extreme ultraviolet (EUV) lithography is regarded as the most promising Next Generation Lithography (NGL) technology. In ITRS 2008 road map, the target of resist performance in EUV lithography is set to be as following. Resolution, line width roughness (LWR) and sensitivity requirements for DRAM half pitch are 16 nm, 0.9 nm 3σ, 10 mJ/cm², respectively.[1] Thus, the requirement of resist performance individually has become very strict demand. Moreover, it becomes the most difficult task to simultaneously fulfill these requirements because there is a trade-off among these requirements.[2] In order to overcome these problems concurrently, it is very important to separately understand reaction mechanisms of the resist pattern formation and inherent resist characteristics such as acid generation efficiency, dissolution kinetics, deprotection reaction and so on.

In pattern formation of chemically amplified resists, the energy is first deposited on resist materials by incident radiation such as EUV radiation. Using the accumulated energy, acids are generated. The acids catalyze pattern formation reactions during PEB via typically the deprotection reaction of polymers, causing the change in alkaline solubility of the resist in exposed areas.[3] It is indispensable for feasibility of EUV lithography to grasp basic chemistry of resist matrices in all stage.

So far, we have estimated acid generation efficiencies in resist matrices and elucidated the reaction mechanism of chemically amplified resist for ionizing radiation such as Electron beam (EB)
and EUV. [4-6, 9] The acid generation mechanism for EUV resists is analogous to that of EB resists. [7-9] However, the absorption of incident EUV photon conforms to Lambert’s Law in EUV resists as well as KrF and ArF photoresists. Thus, it is clarified that EUV resists have some similarity to those in both photoresists and EB resists.

With the reduction in feature sizes, the acid generation efficiency and chemical reactions such as deprotection reaction and dissolution kinetics in resist films becomes critical to accomplish high resolution and high sensitivity. Especially, dissolution kinetics and deprotection reaction of resist films is a key factor to pattern formation. Therefore, it is important to understand dissolution kinetics and deprotection reaction of resist films for the feasibility of EUV lithography.

Because the acids are generated by the ionization of polymer in EUV resists, acid generators hardly affect the absorption coefficient of EUV.[10, 11] Thus, polymers have to be modified in order to adjust the absorption coefficient of EUV resists. Therefore it is essential to investigate polymer structure effects on the dissolution kinetics and deprotection reaction. In this study, polymer structure effects on the dissolution kinetics and deprotection reaction were investigated using a resist development analyzer (RDA-800EUV)[12-13] measurement system (Litho Tech Japan) and deprotection reaction analyzer. [14] The time dependent reactions of the acid-labile protecting group was monitored by FT-IR and the kinetic parameters such as deprotection reaction constants and activation energies were obtained from these data. Also, the simulation of resist profiles for EUV chemically amplified resists was performed by calculation using PROLITH and parameters obtained with EUV exposure tool. Furthermore, the resist patterns were produced by EB and the SEM observation results were compared with the PROLITH simulation results.

2. Experimental

In experiments, PHS derivatives partially-protected by three kinds of protecting groups were used as a polymer (Polymer A–C) as shown in Figure 1. Triphenylsulfonium-triflate (TPS-tf) was used as an acid generator. Propylene glycol monomethyl ether acetate (PGMEA) was used as a casting solvent. Resist film thickness was measured with a film thickness measurement system (Filmetrics F20, spectroscopic ellipsometer) and a Dektak3ST surface profiler.

In the experiments for the analysis of deprotection reaction, resist solutions were spin-coated onto silicon substrates and were pre-baked under the conditions of dissolution kinetics measurements. Then the films were exposed to EUV radiation (Energetic, EQ-10M). The exposed area was approximately 1 × 1 cm². The exposure doses ranged from 0.1 to 1 mJ/cm². After the exposure, they were baked in the temperature range of 80-140 °C for 150 s to estimate deprotection reaction. We used deprotection reaction analyzer to analyze deprotection reactions for EUV lithography. This system makes it possible to measure thin film by using an MCT detector that offered higher detection sensitivity. IR measurements were performed with a resolution of 4 cm⁻¹ over the wave number range from 700 to 4000 cm⁻¹. Typically 64 spectra were averaged from each exposure measurement and single measurement scan was performed with sampling every ca. 0.9 second. By analyzing subtractive spectra between spectrum obtained in resist films and spectrum of the only Si wafer, we can extract information on the resist such as deprotection reaction constant (Kdp), the activation energy of deprotection reactions in resist matrices and so on.

In the sample preparation for dissolution kinetics measurements, resist solutions containing 3.1mol % acid generator were spin-coated onto silicon substrates at 3000 rpm for 30 s to form thin films. After the exposure, dissolution rate measurements were conducted at 110 °C for 60 s under PEB conditions within 1 min of initial exposure to minimize airborne contamination. They were developed in NMD-3 developer [2.38 % tetramethylammonium hydroxide (TMAH) solutions] for 30s at 23 °C and the dissolution kinetics was investigated using a resist development analyzer (RDA) measurement system (Litho Tech Japan). For this analysis tool, the resist dissolution is obtained by measuring the thickness of resist films coated on Si before and after
development. The dissolution rate was estimated by monitoring the variation in interference of a 470 nm wavelength light reflected at the Si substrate and resist film surface as resist film dissolve in TMAH developer.

Also, EUV lithography simulation was performed by using these data obtained with EUV exposure tools. Furthermore, the resist patterns were produced by 75 kV electron beam. The exposure doses ranged from 25 μC/cm² to 50 μC/cm². After the exposure, they were baked in the temperature range of 80-140 °C for 60 s. They were immersion-developed in 2.38 wt% TMAH solutions for 30s and rinsed in water before drying. Resist patterns were recorded with a SEM (JSM-6335F, JEOL).

3. Results and discussion

FT-IR spectroscopy can directly measure changes in the photoactive species by measuring absorbance peaks unique to the photoactive species. Figure 2 shows the representative results of in situ FT-IR measurements of Polymer A during PEB after the exposure to EUV. The x, y and z axis represents the wave number (cm⁻¹), PEB time (s), the absorbance, respectively. The change in the 1156 cm⁻¹ absorption peak of ester (C-O) bonds accompanying deprotection reaction.

FT-IR measurements of Polymer A during PEB after the exposure to EUV. The x, y and z axis represents the wave number (cm⁻¹), PEB time (s), the absorbance, respectively. As shown in Figure 2, the deprotection reactions in polymer A were observed based on the changes in the 1156 cm⁻¹ absorption peak of ester (C-O) bonds accompanying deprotection reaction. With the PEB time elapse, the decrease in the ester bond absorption peak can be seen in Figure 2. Also, we examined the changes in the C-O bond spectra near 1156 cm⁻¹ without the exposure to EUV to investigate whether additional chemical reaction during PEB was occurred. Subtractive spectra weren’t changed in the 1156 cm⁻¹ region. These represent the deprotection reaction proceeds rapidly during PEB after EUV exposure. In the case of Polymer B and C, the changes in the 1230 cm⁻¹ absorption peak of antisymmetric stretching motion (C-O-C) bonds accompanying deprotection reaction and that in the 947 cm⁻¹ absorption peak were observed. However, the changes in the 1230 cm⁻¹ absorption peak was observed during PEB without EUV exposure while it wasn’t observed at the 947 cm⁻¹ absorption peak. So, the change in the 947 cm⁻¹ absorption peak was monitored as previously reported.[15-17]

In order to analyze the kinetics of the deprotection reaction, the change in IR absorbance with exposure time was converted into a normalized protection group concentration. The relative concentration of protected sites was then measured as a function of bake time for many exposure energies and PEB temperatures. By applying a newly deprotection reaction model[18] (equation (1) below) revised with Petersen’s and Ohfuji’s deprotection reaction model[19,20] to obtained results, best fit values for the deprotection reaction rate constant Kdp could be determined.

\[ P = \left(1 + \frac{A E}{1 + \frac{E}{A}} \right) \exp \left( \frac{K_{dp} \cdot t}{E \cdot A} \right) \left[ 1 - \exp \left( -CE \right) \right] \left[ 1 - \exp \left( -K_{loss} \cdot t \right) \right] \]

Where [P] is the relative concentration of protecting group, Kdp is the deprotection reaction constant during PEB (s⁻¹), m is the deprotection reaction order in PEB, E is the exposure energy,
and t is PEB time, $T_2$ is the reaction delay constant in PEB and $\tau_1$ is the average acid lifetime constant in PEB.

Figure 3 shows the relationship between deprotection rate and PEB times, called the deprotection reaction curves, in the temperature range of 80-140 °C for Polymer A, which is protected by 35% protecting groups. The exposure dose was 1.4 mJ/cm². Circles and lines indicate the measured values and the results of fitting to the equation, respectively. As seen in Figure 3, good fits to data were obtained. The deprotection reaction curves in Polymer B and C were also obtained by the same way as polymer A. Based on the deprotection curves shown in Figure 3, deprotection reaction constant (Kdp) and the activation energy of deprotection reactions in Polymer A, B, and C were calculated. Arrhenius plots were made by these Kdp in order to estimate the activation energies. Figure 4 shows Arrhenius plots of Polymer A. As seen Figure 4, Arrhenius plots have different two regions. The Arrhenius plots were composed of different two regions. One is the regions at low temperature where the reaction rate is controlled by the deprotection reaction. The other is that at high temperature where the reaction rate is controlled by acid diffusion. These agreed with the Byers and Petersen model.[24, 25]

The activation energy constants are summarized in Table 1. The variation of Kdp with temperature follows a classical Arrhenius behavior with the activation energy of ca.21.4-27.8 Kcal/mol for polymer A. Though the value obtained was similar to the value reported for DUV, [18, 21] it is less than Ea of 32 kcal/mol value reported by Long et al and J. Hutchinson.[22,23] This might be due to the difference of acid generator used in experimental. For the small protecting groups such as Polymer A and Polymer B, the dependence of activation energy on protecting ratio was small. On the other hand, for bulky protection group, the protecting ratio significantly affected the activation energy probably due to the effect of steric hindrance. Thus, the effect of protecting groups and protecting ratio on activation energy was observed.

Also, the effect of protecting group and protecting ratio on dissolution kinetics was investigated. Figure 5 shows the dissolution rate curve according to protection ratio of Polymer A. The curves relate the resist dissolution rate as a function of exposure dose on a log/log scale. The dissolution parameters such as maximum dissolution rate (Rmax) and minimum dissolution rate (Rmin) were obtained from Figure 5. A slight difference in Rmax was observed due to the difference of resulting products. Also, Rmax at each protecting ratio did not have a significant large difference. On the other hand, Rmin

<table>
<thead>
<tr>
<th>Polymer</th>
<th>(20%)</th>
<th>(25%)</th>
<th>(35%)</th>
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<tr>
<td>Activation energy (kcal/mol)</td>
<td>21.4</td>
<td>38.3</td>
<td>27.8</td>
</tr>
<tr>
<td>Polymer B</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Activation energy (kcal/mol)</td>
<td>3.34</td>
<td>8.64</td>
<td></td>
</tr>
<tr>
<td>Polymer C</td>
<td>(10%)</td>
<td>(20%)</td>
<td>(25%)</td>
</tr>
<tr>
<td>Activation energy (kcal/mol)</td>
<td>1.16</td>
<td>1.16</td>
<td>3.20</td>
</tr>
</tbody>
</table>
decreased with increase of protecting ratio. Higher protect ratio provided increased inhibition. A large value of Rmin signifies that the resist easily and smoothly dissolves into alkaline developer. These results suggest that a higher protecting ratio increase the resists potential for higher resolution capability as previously reported. [26, 30] It is reported that the dissolution of resists to an alkaline developer is attributed to the penetration of base molecules from the developer into the resist by base molecule diffusion through the hydrophilic sites of the polymers. This is called as the concept of the “penetration zone”. [31-33] The speed of penetration increases with the increasing volume of the hydrophilic space of the resist film. On the other hand, because hydroxy group of base resin is hydrophilic TMAH solution, it can easily penetrate into resist film as reported. [27, 28, 34] Thus, it is supposed that the hydrophilic substituent content of the base polymer determines the degree of penetrability of the base molecules and the dissolution rate of the resist to the developer. The speed of TMAH penetration into each resist film is different for each resist in our study. This indicated that the dominant rate-determining step in these resists is TMAH penetration into the resist film.[35] The dissolution rates was affected by changing the protection ratio of polymer.

Furthermore, we performed EUV lithography simulation by using these data obtained with EUV exposure tools. The calculation parameters of Polymer on PROLITH are shown in Table 2. The partially coherence factor of the illumination system was 0.5. The resist line width was set to 26 nm and 50 nm 1:1 line& space pattern. The exposure dose is set to 20, 25, 30 mJ/cm². At the film thickness of 50 nm, an almost vertical is obtained for Polymer A. On the other hands, at the film thickness of 26 nm there is considerable bridge of pattern side walls, and good quality patterns were not obtained as shown in Figure 6.

Table 2. Calculation parameters of Polymer A on PROLITH.

<table>
<thead>
<tr>
<th>Development Parameters</th>
<th>Development Rmax</th>
<th>Development Rmin</th>
<th>Development Mth</th>
<th>Development n</th>
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<tr>
<td></td>
<td>46.6 mm/s</td>
<td>0.10 mm/s</td>
<td>0.39</td>
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<td>PER Parameters</td>
<td>Thermal Decomp. Ea</td>
<td>27.8 kJ/mol</td>
<td>Thermal Decomp. Int/Ar</td>
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<tr>
<td>ABC Parameters</td>
<td>A</td>
<td>0 [gm]</td>
<td>B</td>
<td>1.06 [gm]</td>
</tr>
<tr>
<td></td>
<td>C</td>
<td>0.0097 [cm/mJ]</td>
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<td></td>
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Fig. 6 50 nm and 26 nm L&S pattern profiles of Polymer A resist, calculated by PROLITH.

Fig. 7 SEM image of L&S pattern in Polymer A resist by using 75 kV EB.

The resist patterns were produced by EB and SEM observation result was compared with PROLITH simulation result. Figure 7 shows representative SEM images of Polymer A. As for the resolution, the resolution limit by using EB is almost same as prediction based on PROLITH. PROLITH might suggest the possibility of rough examination of resolution limits for EUV resist.

4. Conclusion

Polymer structure effects on the dissolution kinetics and deprotection reaction were investigated to understand inherent EUV resist characteristics. The effect of protecting ratio on activation energy was observed. For small protecting group, dependence of activation energy on protecting ratio was small. On the other hands, for bulky protecting group protecting ratio significantly affect the activation energy probably due to the effect of steric hindrance. Also, the deterioration of resist sensitivity was observed with increase of protecting ratio while the dissolution slope increases with increase of protecting ratio. Thus, the dissolution rates and sensitivity were more affected by changing the protection ratio of polymer than the type of protecting group. Also, we performed EUV lithography simulation by using PROLITH. It is important for EUV lithography to select appropriate protecting group and protecting ratio.
Acknowledgement

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