Underlayer designs to enhance the performance of EUV resists

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ABSTRACT

Extreme ultraviolet (EUV) lithography has gained momentum as the method of choice for <32-nm half-pitch device fabrication. In this paper, we describe our initial attempts to increase an EUV resist’s sensitivity without compromising resolution and line roughness via introduction of a thermally crosslinkable underlayer. The main purpose is to test the possibility of using a combination of photoacid generators (PAGs) and EUV sensitizers (phenol type) in the underlayer designs to enhance the overall performance of EUV resists. We have demonstrated the possible benefits of adding an EUV underlayer into the regular EUV litho stack and investigated the effect of PAG types and loadings on the photospeed and litho performance of three different EUV resists.

Keywords: EUV, underlayer, lithography, photospeed, resolution, LWR, RLS tradeoff, photoacid generator, PAG

1. INTRODUCTION

EUV lithography (EUVL) is expected to offer a single-exposure solution for use in manufacturing at 22-nm half pitch, and possibly for some applications at 32-nm half pitch.\textsuperscript{1,2} However, in order to successfully implement EUVL in time, several technical hurdles must be overcome including: mask fabrication with low defects\textsuperscript{3}; reliable EUV sources with high output power and sufficient lifetime\textsuperscript{2}; contamination control of all mirrors\textsuperscript{5}; high throughput; photoresists with sufficiently low line width roughness (LWR) and low exposure dose. For resist requirements, the international technology roadmap for semiconductors (ITRS)\textsuperscript{6} set targets for 3-σ LWR at < 3 nm and for sensitivity at < 10 mJ/cm\textsuperscript{2}, which, to date, has not been achieved simultaneously in an EUV resist. Earlier research has actually found that most efforts simply result in a compromise between LWR, resolution, and sensitivity, e.g., good sensitivity but rough lines, or improved LWR but poor resolution. Although the design of new EUV photoresists and an improved understanding of the patterning mechanism has allowed significant progress for simultaneously achieving improved EUV resolution, LWR, and sensitivity (RLS), the trade-off is clearly a natural limitation for chemically amplified resists.\textsuperscript{7,8,9}

In previous studies and publications, we have demonstrated the benefits of introducing an EUV underlayer beneath the resist. Specifically, we provided initial data on the relationship between film density/adsorption and EUV litho performance.\textsuperscript{10} Additional benefits of using an EUV underlayer include, but are not limited to, planarization of a substrate with topography, protection of underlying layers from possible damage caused by high-energy EUV photons, improved adhesion for high-aspect-ratio resist profiles, and improved etch selectivity for better pattern transfer. Our goal in this paper is to use a spin-on EUV underlayer beneath the resist to harvest the extra energy from EUV photons. The energy is then redirected back to the resist in the form of photons, secondary electrons, or generated photoacids to enhance the resist’s photosensitivity.

The underlayer is a crosslinkable polymeric system that can be coated onto a substrate through a regular spin-on process and then thermally cured so that the photoresist can be applied thereafter. Most underlayers have exhibited less outgassing than positive tone photoresists during EUV exposure.\textsuperscript{11} The polymer was chemically modified to have relatively high EUV adsorption and functional additives were included in the underlayer formulations to enhance the energy harvest and transfer process. In this paper, we will describe some of our efforts on various underlayer designs and their impacts on resist performance, mainly focusing on the possibility of using the combination of photoacid generators (PAGs) and EUV sensitizers (phenol type) in the underlayer to enhance EUV resist RLS performance.
2. EXPERIMENT

2.1 Material and Formulation

All the underlayers studied in this paper are based on a crosslinkable linear polymer system. As shown in Figure 1 below, there are two basic polymer platforms involved in this study: a methacrylate platform A and polyhydroxy styrene (PHS) platform B. In platform A, various PAGs, including triphenylsulfonium triflate (TPS-OTf), TPS nonaflate (TPS-ONf), and PAG-C1, were physically blended into the formulations to produce UL2, UL3, and UL4, respectively. In the case of UL4, different amounts of PAG-C1 were added to generate a series of UL4-10 (10 wt%), UL4-16 (16 wt%), and UL4-20 (20 wt%). 2,5-dimethyl phenol (2,5-DiMe Ph) was added as an EUV sensitizer together with TPS-OTf to obtain UL5. Platform B bears triflate PAG and phenoxy groups on the polymer backbone, so no additional ingredients were needed to produce UL6. The polymer originally features 30% of PHS functionalized as a triflate, so the initial formulation using this polymer and crosslinker was named UL6-30. For comparison, PHS with no triflate moieties was blended in at one-half and two times the weight of this triflate functionalized PHS to make UL6-20 and UL6-10, which means 20% or 10% of the total PHS units were triflated, respectively.

![Chemical structures of two polymer platforms and six different underlayer formulations tested as part of this study.](figure1.png)

<table>
<thead>
<tr>
<th>Underlayers</th>
<th>Platform</th>
<th>PAG</th>
<th>EUV sensitizer</th>
</tr>
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<tbody>
<tr>
<td>UL1</td>
<td>A</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>UL2</td>
<td>A</td>
<td>TPS-OTf</td>
<td>None</td>
</tr>
<tr>
<td>UL3</td>
<td>A</td>
<td>TPS-ONf</td>
<td>None</td>
</tr>
<tr>
<td>UL4 (-10, -16, -20)</td>
<td>A</td>
<td>PAG-C1</td>
<td>None</td>
</tr>
<tr>
<td>UL5</td>
<td>A</td>
<td>TPS-OTf</td>
<td>2,5-DiMe Phenol</td>
</tr>
<tr>
<td>UL6 (-10, -20, -30)</td>
<td>B</td>
<td>None</td>
<td>None</td>
</tr>
</tbody>
</table>

Figure 1. The chemical structures of two polymer platforms and six different underlayer formulations tested as part of this study.

2.2 Process Condition and Measurement

Polymer, crosslinker, and additives (if any) were dissolved completely in typical semiconductor-grade solvents (such as PGME, PGMEA, and combinations of the two) and filtered through a 0.1-μm filter before use. Each of the final
formulations (TSC = 1.5 ~ 2.5%) was spin-coated at 1500-3000 rpm onto a silicon substrate and then baked at 160°-
205°C for 60 seconds. The resultant film thicknesses ranged from 20 to 60 nm, depending on the specific requirement of
the lithography stack. The film was rinsed with ethyl lactate to test the film’s solvent resistance (EL strip) and the film
was also immersed in photoresist developer (TMAH base solutions) without light exposure to evaluate the dark loss. All
the formulations used here were under 5% EL stripping and dark loss without specification.

2.3 Instrumentation

2.3.1 EUV Exposure and TOP CD SEM

EUV exposures were performed on a microexposure tool (MET) either at the Advanced Light Source (ALS) in Berkeley
or at Intel in Hillsboro, OR. The ALS-MET utilizes a 5x-reduction, 0.3-NA optic, providing a 200×600 μm exposure
field at the wafer plane. All processing was performed using Berkeley BKMs as previously reported.13 The Intel-MET
is a micro-field 600x600 mm², 0.3 numerical aperture (NA), two mirror lithography exposure tool, made by Exitech, Ltd.
with resolution in excess of 30nm. The illumination settings used were annular with s inner=0.36 and s outer=0.55,
normalized with respect to the pupil radius of 1. All processing was done on 300mm underlayer-coated Si wafers using
a linked TEL-ACT12 track in Ronler Processing 1 (RP1), a Class 1 clean room fab. The CDs and LWR measurements
were made using a CD Scanning Electron Microscope (SEM). The methodology used to measure LWR has been
detailed previously.14 All of the LWR values reported in this article are an average of at least five to seven lines repeated
at five different locations in the array. All photoresist testing was done using a 60nm post-apply bake film thickness in
order to maintain an aspect ratio of roughly 2:1 to help prevent pattern collapse. An aspect ratio of 2:1 balances the
opposing needs to minimize resist thickness to reduce pattern collapse versus maximizing the resist thickness to help
ensure that film absorbance is high enough such that sufficient aerial image contrast and photon absorption are achieved.

2.3.2 Cross-section SEM

The wafers were then sent to CeriumLabs in Austin, TX, for cross-section and imaging. The cross-section images shown
in Figure 5 are after deposition with metals by the e-beam in the focused ion beam (FIB) and imaged using Hitachi
S4800 high-resolution SEM/STEM.

2.3.3 Time-of-flight Secondary-ion Mass Spectrometry (ToF-SIMS)

ToF-SIMS is a surface-sensitive spectroscopy that uses a pulsed ion beam (Cs or microfocused Ga) to remove molecules
from the very outermost surface of the sample. The particles are removed from atomic monolayers on the surface
(secondary ions). These particles are then accelerated into a "flight tube" and their mass is determined by measuring the
exact time at which they reach the detector (i.e. time-of-flight). Depth profiles are produced by removal of surface layers
by sputtering under the ion beam, which is used as the technique to probe the PAG distribution along the depth of the
resist and underlayer.

3. RESULTS AND DISCUSSION

3.1 The Effect of Adding Underlayers Beneath an EUV Resist

It has been slowly recognized that a properly selected underlayer between substrate and EUV resist could contribute in
enhancing the overall EUV resists’ performance. UL1 is one of the first-generation underlayers, and we used it here as a
baseline for studies described in this paper. As clearly shown in Figure 2, under the same tool setting, exposure
condition, and photoresist, there is an obvious difference between the performance of TOK-P1123 coated directly on
HMDS-primed silicon wafers and when coated onto UL1. The roughness of 40- and 32-nm half-pitch (HP) lines can be
significantly decreased with an EUV underlayer present. When tested with Resist C at Intel, the LWR was also slightly
reduced for both 40- and 30-nm HP lines by using UL1. However, at 32-nm HP the semi-isolated P1123 resist lines
started to collapse on UL1, and the dose required for 30-nm HP Resist C patterning increased. Therefore, adding an EUV
underlayer shows a trend of slightly reducing the line roughness, but may be subject to pattern collapsing and / or
increased dose to size at 30-nm HP, depending on the EUV resists utilized. The explanation for LWR improvement by
using an UL is not clear at this point, and is beyond the scope of this paper. The improvement of the adhesion to prevent
pattern collapsing is under investigation now. Our main goal here is to understand, using UL1 as a control, if the addition of PAGs and/or EUV sensitizer can increase the photosensitivity of the resist without negatively affecting LWR or resolution.

Figure 2. The comparison of top-down SEM images, at 40-/32-nm, and 40-/30-nm between HMDS-primed wafers and UL1, using either TOK-P1123 at ALS-MET Berkeley or Resist C at Intel-MET.

3.2 The Photospeed Enhancement of EUV Resist by Adding PAGs and/or EUV Sensitizers into the Underlayer

In an effort to increase the photospeed of currently available EUV photoresist materials, we have investigated various additives to the baseline UL1. To increase resist photospeed, the key is to harvest as much energy as possible and then feed it back to the resist in different forms, e.g., photons, secondary electrons, or photo-generated acids. The first study was carried out by adding PAGs and energy sensitizers into the underlayer design in an effort to enhance the photoresist’s performance, i.e., increased photospeed without compromising LWR and/or ultimate resolution.

Three different PAGs - TPS-OTf, TPS-ONf, and PAG-C1 - were added to UL1 at the same molar concentration. The same EUV resist (TOK-P1123) and illumination conditions were utilized in this study. Figure 3 shows the contrast curves for five different substrates, primed silicon substrate (HMDS), UL1, and UL1 doped with TPS-OTf (UL2), TPS-ONf (UL3), and PAG-C1 (UL4). Dose to clear (E0) is defined as the dose at the point when film thickness (FT) decreases to < 1% of the original FT or when the curve slope starts to approach zero. The contrast (in the unit = mJ⁻¹) was also calculated as the maximum slope of each contrast curve. Clearly, the addition of PAGs into the UL decreased the E0 significantly, by 16% in the case of the C1 PAG. This is not too surprising because PAGs activated during exposure will generate photoacid that can help to clear the resist. An additional observation is that the C1 PAG has the largest effect while triflate shows the smallest, when added at the same molar concentration. The calculated slopes/contrasts demonstrated the same trend with increased contrast from HMDS-primed wafer, to UL1, UL2, UL3, and UL4. We tentatively attribute this correlation to either acidic strength caused by various anions, or the different migration length/depth into the top resist due to the inherent difference in molecular structure. However, more experiments are needed to reach a definite conclusion.
Adding EUV sensitizers (phenol types) did not show effects as that were as obvious as the addition of PAGs. We explored different types of phenolic sensitizers, including but not limited to 2,5-dimethyl phenol, 4-nitro-phenol, 4-methoxy-phenol, 2,6-dinitro-4-bromo-phenol, and various bisphenol and polymer bounded phenols. Some of the phenols, such as 2,5-demethyl phenol, have shown great effect in enhancing photospeed when added directly into photoresist. Unfortunately, only a slight differences in $E_0$ with and without EUV sensitizers (phenols) in the underlayer design were observed, and no clear trends were present among different types of phenols at various concentrations. Part of the explanation might be due to the fact that most phenol molecules readily sublime, such that few are left in the matrix after our PAB. We also believe that the EUV sensitizers need to be in the proximity of the PAG to really see an effect, which is challenging for blended materials because of inhomogeneities and the high crosslinking density in UL1-UL4. We will address this challenge in a later section of the paper.

3.3 The General Effect of Added PAG and Sensitizer on Pattern Profiles

To check the effect on line roughness and ultimate resolution, wafers were patterned using the ALS-MET at Berkeley with TOK-P1123. A 5 X 9 FEM was used, with 100-nm focus step and 7% of central dose as dose step. Figure 4 shows the side-by-side comparison of 32- to 60-nm half-pitch top CD SEMs from UL1, UL2 (with TPS-OTf), and UL5 (with both TPS-OTf and 2,5-dimethyl phenol). There is some decrease in $E_{size}$ with added PAG and/ or phenols, 7%-14% varied at different pitches. However, the imaging quality deteriorated with the addition of PAG and/ or phenol. Especially at smaller pitches, the line roughness increased rapidly (7%-26%) and lines started to fall off at 32- and 34-nm HP. Clearly, the LWR and resolution were more compromised than the modest improvement in photospeed.

To more quantitatively analyze the data to examine the RLS trade-off, the concept of Z factor was used to evaluate the balance of resolution, LER/ LWR, and photospeed. Z factor, as described elsewhere, is defined as:

$$Z \text{ factor} = (\text{resolution})^3 \times (\text{min LER})^2 \times E_{size}$$

The current target for 32-nm HP line/ space patterning is 2.0-nm 3-σ LWR (or 1.4-nm LER) and 10 mJ/cm² dose to size, or $Z$-factor (32-nm target) = 6.6E-09, which is defined as the target. A normalized value can then defined as $nZ$ (32 nm),

$$nZ_{32} = Z \text{ factor} / Z \text{ factor (32-nm target)}$$

According to the Top SEM shown in Figure 4, the $nZ_{32}$ calculated for each underlayer at the ultimate resolution are: 14.1 for UL1 at 34 nm HP, 22.3 for UL2 at 36 nm HP, and 17.4 for UL3 at 36 nm HP. Clearly there is no significant RLS improvement observed, although UL5 (PAG in combination with sensitizer) showed slightly better performance than UL2 (PAG only). (Table 1)
Cross-section SEM images were taken at the best dose and focus for 36-, 40-, and 60-nm HP to evaluate the pattern profile and line shape. As shown in Figure 5, footing was observed with UL1, and became more severe as CD size decreased from 60 nm to 40 nm to 36 nm. Adding PAG and/ or phenols not only decreases the required dose to size, which is in agreement with top CD SEM observations, but also helps to keep the resist’s profile straight by reducing footing and scumming. To investigate the cause of footing and how a PAG in the UL formulation helps, TOF-SIMS was performed to track fluorine (F)/ PAG distribution along film depth to the substrate, as all three PAGs used here bear the distinctive F atom compared to the rest of the hydrocarbon-based resists/ ULs. The SIMS data first confirmed the composition of these four different ULs: no PAG in UL1, and different PAGs in UL2, UL3, and UL4 with equal molar concentrations. Because triflate has only 3 F atoms per molecule while both nonaflate and C1 PAG have 9 F atoms per molecule, it is not surprising that the F peak in UL2 is much lower than in UL3 and UL4. It is also clear that the distribution of PAG inside the resist is not uniform, with a deficient area near the photoresist / UL interface, which at least partially explains the formation of footing. Therefore, the photoacids generated and then diffused from the PAGs of the top UL can balance this deficiency, and keep the resist profile straighter. However, untamed PAG/ photoacid diffusion from the UL can also causes line collapse, especially semi-isolated lines, which explains why the ultimate resolution was not as good as the baseline UL1, where no PAG was added.
**Figure 5.** Cross-section SEM images of 60-, 40-, and 36-nm HP lines on UL1, UL2, and UL5.

**Figure 6.** SIMS profile of TOK-P1123 (70 nm) resist on UL1, UL2, UL3, and UL4 (40 nm).
3.4 The Effect of PAG Amount on Dose to Size of Resist

To examine the effect of different PAG loadings, various amounts of PAG-C1 were used in the formulation of UL4 at the levels of 10-wt%, 16-wt%, and 20-wt%. The resulting formulations were called UL4-10, UL4-16, and UL4-20, respectively. In this study, all underlayers were coated at 60 nm FT using Resist C. Dose to clear (E₀) and dose to size at 30-nm HP (E₃₀) were evaluated with Intel’s MET tool. First of all, a 24% E₀ decrease was observed, which is in agreement with previous results at ALS LBL using TOK-P1123 resist. Clearly the increased photospeed resulting from the addition of PAG to the UL was independent of the exposure tool and EUV resist used. Second, a 4% E₃₀ decrease further confirmed that the photospeed increase is not only a macroscopic phenomenon but also happens at the microscopic scale. Another important trend is that increasing PAG loading does not generate a greater photospeed increase; rather both E₀ and E₃₀ saturated at a certain value after 10-wt% PAG additions. From the PAG distribution indicated in Figure 6, the possible explanation might be that increased PAG loading will only increase the peak height (central concentration) inside the UL and not increase the number of photoacids that can actually diffuse into the top resist. The real PAGs that contribute lie only at the interface, and their concentrations are not significantly effected by the total PAG concentration inside the UL. This is a very valuable observation for future investigative direction: focusing on the increase of local PAG concentration at the resist/UL interface.

<table>
<thead>
<tr>
<th></th>
<th>UL1</th>
<th>UL4-10</th>
<th>UL4-16</th>
<th>UL4-20</th>
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<tr>
<td>Dose to clear (E₀)</td>
<td>6.25</td>
<td>4.75</td>
<td>4.75</td>
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<tr>
<td>Dose to 30nm (E₃₀)</td>
<td>12.25</td>
<td>11.75</td>
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</table>

Table 2. The dose to clear and dose to size 30-nm HP (in mJ/cm²) of Resist C on UL1, UL4-10, UL4-16, and UL4-20.

3.5 PAG Comparison Between Triflate and PAG C1

In an effort to utilize the PAG in a more controlled way to maintain the benefit of decreased Eₚsize without sacrificing LWR and resolution, a PAG utilizing a bulkier anion was employed. TPS-triflate has been known to have a long diffusion length that can lead to unwanted line collapse and lifting. PAG C1 with a branched multifluorocarbon functionalized anion, was used in the following study in comparison with TPS-triflate to see if the selection of PAG would lead to a difference in patterning quality. Wafers were patterned using the Intel-MET with Resist C. An 11 X 21 FEM was used, with 40-nm focus step and 0.75-mJ/cm² dose step.

The results are shown in Figure 7, comparing the Eₚsize, LWR, Z-factor, and depth of focus (DOF) for UL1, UL2 (triflate), and UL4 (PAG C1). Because of the different resist and tool used this time, the comparison between UL1 and UL2 is slightly different from what was observed in Figure 4. However, there is still no convincing increase in Z factor from the addition of TPS-triflate to the UL. Focusing on the difference between triflate and PAG C1, it is easy to notice that UL4 with PAG C1 did generate an obvious advantage over UL2 with triflate PAG – smoother lines, smaller Z-factor, and bigger DOF for the same Eₚsize. This actually confirmed our initial assumption that by using an equal amount of PAG C1 instead of triflate, the same photospeed increase could be achieved while not greatly impacting LWR and resolution due to a more controllable diffusion length of a bulky anion.
3.6 An Improved System – Platform B and UL6 Series

Through the initial studies on the Platform A–based system, we found that, due to uncontrolled photoacid diffusion, the benefit we gained in photospeed increase was compromised by the loss of resolution and an increase in line edge roughness. In addition, it appears that in order to make the EUV sensitizer work together with PAG, they need to be put in close proximity to each other. Finally, physically blending small molecules in the underlayer formulation does not seem very efficient and is hard to control. Based on these three observations, we designed a new polymer system that includes a crosslinkable polymer matrix, a phenolic-type sensitizer anchored to the polymer backbone, and a “cation-bound” covalently attached PAG, shown as Platform B in Figure 1. With such a design, first, the PAG is bound to the polymer so that it is more evenly distributed in the underlayer and less likely to leach into the photoresist during resist coating; second, the phenol is part of the polymer such that it is not as apt to sublime as most phenol-type small molecules; and third, the PAG and phenol are held in proximity to each other for more efficient energy transfer.

In this study, an open-source resist – TER60AB – was used, which comprises a 60% hydroxystyrene, 20% styrene, 20% t-butyl acrylate terpolymer, TPS-triflate as PAG, and trioctylamine as a base quencher. Although it is not the best-performing EUV resist, its open-source structure and composition gave us the opportunity to understand how the resist and underlayer interact together.

During the initial tests, it was found that UL6-10 gave the best performance, as shown in Figure 8. Due to a dose shift during the exposure slot, there is no exact dose information associated with each SEM, but through a reasonable estimation, we believe the maximum dose would be ~ 9 mJ/cm². If this assumption is correct, the SEM of 32-nm HP shown below is much better than the 32-nm lines illustrated in Figure 4, i.e., the lines are smoother with reduced pattern collapse. The Z factor calculated is 6.8E-08 and nZ32 = 10.3, much closer to the 32-nm HP target. In addition, the ultimate resolution extended to 28-nm HP (with 26.8-nm actual CD). By calculation, the Z factor is 5.8E-08 and nZ32 = 8.9. With an open-source triflate-based resist, this result is very encouraging, and we are in the process of improving the structure design and formulation.

Figure 7. The comparison of Resist C performance on UL1, UL2, and UL4-16.
4. SUMMARY

In summary, to test the idea of improving EUV resists’ performance by introducing an EUV underlayer with specially designed additives, six types (10 actual formulations) of underlayers were investigated with three different resists (TOK-P1123, Resist C, and open-source TER60AB) on MET EUV exposure tools at both Berkeley and Intel. The use of an appropriate underlayer can clearly impose some benefits on the overall EUVL performance (e.g. slightly decreased LWR), independent of the resist and tool used. The addition of PAG into an EUV underlayer reproducibly showed the ability to increase the photospeed of the resists, i.e., 7%-25% decrease of $E_0$ and $E_{size}$. Underlayers with PAG (or together with sensitizer) can also reduce footing and scumming between patterned lines, keeping the resist shape more square and straight. However, untamed photoacid diffusion still causes this improvement to be subject to the RLS trade-off by compromising LER and resolution. Therefore, we have not observed a significant RLS gain using the underlayers reported herein. Recent designs that incorporate PAG and possible EUV sensitizers into the polymer backbone provided very promising initial results. Further investigation is underway and will be reported in the future.

5. ACKNOWLEDGEMENT

The authors would like to thank Huiying Liu (Intel) for TOF-SIMS analysis and JSR Corporation / JSR Micro, Inc. for donation of the TER60 resist employed in this study.

6. REFERENCES


Figure 8. SEM images of open-source resist TER60AB on UL6-10 at 40-, 32-, and 28-nm HP lines.

CD = 45.8 nm CD = 31.3 nm CD = 26.8 nm
LWR = 8.9 nm LWR = 7.0 nm LWR = 8.2 nm