Nanoparticle Contamination Control and Metrology for the EUVL Systems

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Outline

- Background and Motivation
- Protection Schemes for EUVL Masks
 - Carriers at Atmospheric Pressure
 - Scanners at below 100 mTorr
- Nanoparticle Metrology and AMC Issues
 - Standardization of Nanoparticles
 - Mask Deposition and AMC Issues



Background and Motivation



- Pellicles are unavailable for protecting the EUVL masks due to high absorption of EUV beam in most solid materials
- EUVL masks need to be protected against all particles > about 20 nm



Protection Schemes

The Intel project started in 2004. Particle contamination of EUVL photomasks was unknown. It was feared that thousands of particles might deposit on the mask during each operation. We need to investigate a broad range of protection schemes.

- Mask inside a carrier or scanner
- Cover plate to reduce risk volume
- Critical surface upside down to avoid gravitational settling (Cover plate underneath mask during shipping, storage, and pump down)
- Electric field to make use of electrophoresis
- Thermal gradient to make use
 of thermophoresis



 Particle trap surrounding mask to avoid particle penetration from the side



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Particle Source Identification

ATOFMS

Mask Scan





 Complex organic compound or mixture -possibly polymer Contact points between the mask surface and pins

Particles come mostly from contact points between mask surface and pins

Yook et al., IEEE Trans. Semi. Manu. 20(2): 176-186 (2007).



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Validation of Pozzetta Carrier Design on Particle **Generation during Real Shipping**



• The standoff-support generates almost no particles.



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Study of Various Protection Schemes inside a Carrier



Effect of Cover Plate Protection ($d_p = 10$ nm)



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Thermophoresis Test Set Up

No Gradient

With Gradient



Vacuum chamber











Thermophoresis at 100 mTorr, 2 cm Gap



Simulations at 50 mTorr 125 nm, 1 cm Gap, $v_i = 6.5$ m/s



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New NIST Nanoparticle Standards: 60 nm and 100 nm SRM

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Measurement of 100 nm and 60 nm Particle Standards by Differential Mobility Analysis

Number 4

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george mulholland@nist.gov michelle.donnelly@nist.gov charles hagwood@nist.gov vince.hackley@nist.gov The peak particle size and expanded uncertainties (95 % confidence interval) for two new particle calibration standards are measured as 101.8 nm ± 1.1 nm and $60.39 \text{ nm} \pm 0.63 \text{ nm}$. The particle samples are polystyrene spheres suspended in filtered, deionized water at a mass fraction of about 0.5 %. The size distribution measurements of aerosolized particles are made using a differential mobility analyzer (DMA) system calibrated using SRM® 1963 (100.7 nm polystyrene spheres). An electrospray zerosol generator was used for generating the 60 nm aerosol to almost eliminate the generation of multiply charged dimers and trimers and to minimize the effect of non-volatile contaminants increasing the particle size. The testing for the homogeneity of the samples and for the presence of multimers using dynamic light scattering is described. The use of the transfer function integral in the calibration of the DMA is shown to reduce the uncertainty in the measurement of the peak particle size compared to the approach based on the peak in the concentration vs. voltage distribution. A modified aerosol/sheath inlet, recirculating sheath flow, a high ratio of sheath flow to the aerosol flow, and accurate pressure, temperature, and voltage measurements have increased the resolution and accuracy of the measurements. A significant

consideration in the uncertainty analysis

July-August 2006

was the correlation between the slip correction of the calibration particle and the measured particle. Including the correlation reduced the expanded uncertainty from approximately 1.8 % of the particle size to about 1.0 %. The effect of non-volatile contaminants in the polystyrene suspensions on the peak particle size and the uncertainty in the size is determined. The full size distributions for both the 60 nm and 100 nm spheres are tabulated and selected mean sizes including the number mean diameter and the dynamic light scattering mean diameter are computed. The use of these particles for calibrating DMAs and for making deposition standards to be used with surface scanning inspection systems is discussed.

Key words: differential mobility analysis; dynamic light contering; electrical mobility; electrospray aerosol generation; particle size calibration standards; transfer function.

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Nanometer Differential Mobility Analyzer (Nano-DMA)

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National Institute of Standards & Technology

Certificate Standard Reference Material 1963

Nominal 0.1 µm Diameter Polystyrene Spheres

This Standard Reference Material (SRM) is intended primarily for use as a primary reference standard for the calibration of particle size measuring instruments including optical and electron microscopes. The SRM consists of 5 mL of carborylated polystyrone spheres in water as a weight concentration of about 0.5%. It is supplied in a discensing vial.

The number average particle diameter was measured in air as an aerosol by electrical mobility measurements. The certified value is:

Number Average Diameter, µm

0.3007

± 0.0020

Uncertainty, am

The uncertainty includes both random and systematic errors. The total random uncertainty is $0.00055 \,\mu$ m (99% confidence interval), and a conservative estimate of the systematic error is $0.0014 \,\mu$ m.

The size distribution of the polyhymne spheros, as determined by electrical mobility measurements, is anreas with a standard deviation of 0.0018 μ m excluding outliers. The number of undersized particles is negligible and the number of overviewed particles (diameters greater than 0.2 μ m) is less than 0.1%.



Issues with PSL Particle Standard

- Different light scattering than particles from processes
- Decomposition from exposure to deep ultra-violet (DUV) lights
- Deformation due to adhesion forces



A 1.3 μm PSL sphere after adhering to a chromium surface for 24 hrs. From Dahneke, B. "The influence of flattening on the adhesion of particles," J. Colloid Interface Sci., vol. 40(1), (1972).



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Standard Particle Deposition for Scanner Calibration

- Calibration of surface inspection tools with particles of different materials
- Development of accurate size standards
- Providing samples for cleaning studies



Haze Observed under Atmospheric and Vacuum Conditions

50nm SiO₂. Target deposition area: 1inch spot size at the center. Testingtime: 2 min. (Atmospheric Pressure)

100 nm PSL particle. (Main Chamber p = 50 mTorr). Testing time: 1.5 hours







Airborne Molecular Contaminants (AMCs) Classification of AMCs



Controlled Particle Deposition on Mask Blanks

Deposition Plan

(~ 2000 particles)



Mask Scan Area (140 mm x 140 mm)

- Known material
- Known number of particles
- NIST-traceable particle size
- Controlled deposition spot size

Detection of Particles

on a Quartz Mask





Summary

- Experimental methods and models have been developed to evaluate protection schemes for masks in carrier or vacuum tools.
- New carriers with tapered standoff generates almost no particles during shipping.
- Face-down mounting and cover plate are very effective in protection.
- Thermophoresis is most helpful to protect against particles driven by diffusion.
- Method has been developed to deposit standard nanoparticles for inspection tool calibration.
- Method has been developed to avoid haze formation caused by AMC.



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