

Proximity Effect in E-Beam Lithography

Overview and Agenda

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BEAMER

LAB

TRACER

Pro SEM

MASKER

Part	Subject	Date
1	Electron Scattering and Proximity Effect	07-Oct-2020, 6:00pm CEST, 12:00pm EDT, 9:00am PDT
2	Dose PEC Algorithm and Parameter	14-Oct-2020, 6:00pm CEST, 12:00pm EDT, 9:00am PDT
3	Optimization of Dose PEC Parameter	21-Oct-2020, 6:00pm CEST, 12:00pm EDT, 9:00am PDT
4	Process Effect, Calibration and Correction	28-Oct-2020, 5:00pm CET, 12:00pm EDT, 9:00am PDT
5	Shape PEC – “ODUS” Contrast Enhancement	04-Nov-2020, 6:00pm CET, 12:00pm EST, 9:00am PST
	Break	11-Nov-2020 -- No Session
6	3D Surface PEC for greyscale lithography	18-Nov-2020, 6:00pm CET, 12:00pm EST, 9:00am PST
	Thanksgiving Week	25-Nov-2020 -- No Session
7	T-Gate PEC	02-Dec-2020, 6:00pm CET, 12:00pm EST, 9:00am PST

- The webinar series will explain one of the most important techniques in advanced e-beam lithography. Modern E-beam systems are able to form small spot sizes in nm range. In principle this enables to achieve feature sizes in nm-range. In practice this is limited by physics, chemistry and tool limitations...

Proximity Effect in E-Beam Lithography

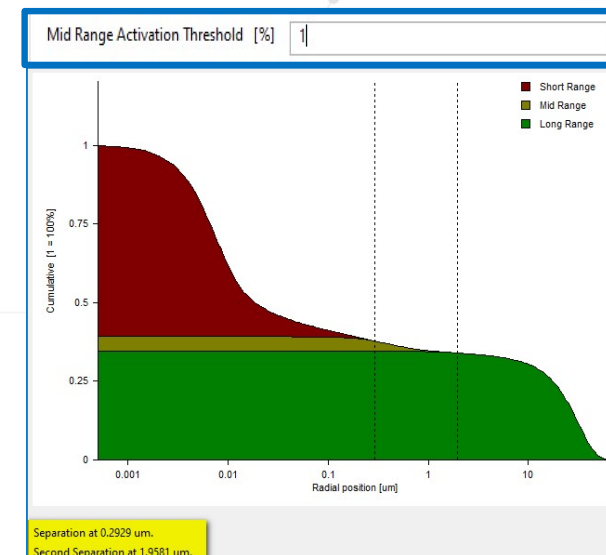
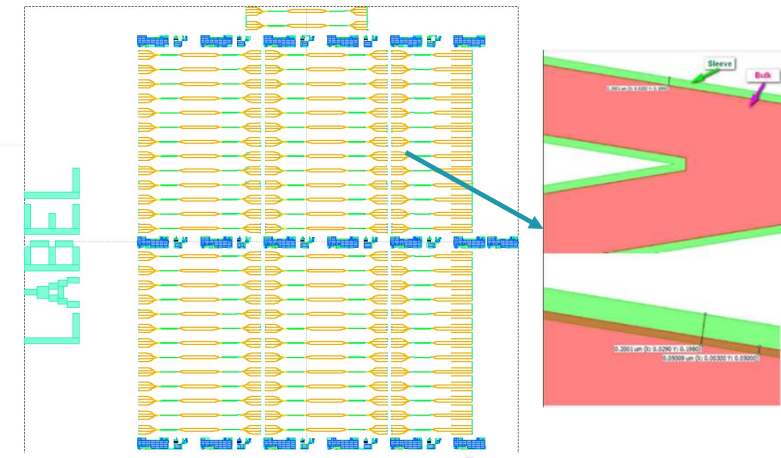
Part 4: Process Effect
Calibration and Correction



- Part 3 Summary: Dose PEC Parameters
- Process Effects and Major Parameters
- Calibration Procedure
- Advanced Model Parameters
- Summary
- Q&A

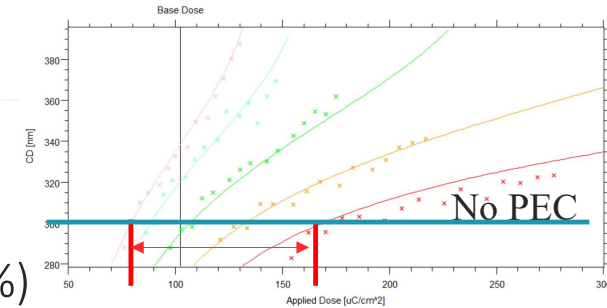
General Dose PEC Parameters

- Include all (and only) pattern to be exposed into the resist
 - PEC does maintain layers (e.g. for bulk-sleeve, writing order control)
 - Include non-critical layer in PEC, but exclude LR fracturing assigns on dose to the feature and considers energy contribution
 - Non critical out of influence range (pads, label,) may be excluded
- PEC can be only as good as the correction function (PSF)
 - Monte-Carlo (table defined) PSF is preferred
 - Including Short Range correction requires “Effective Short Range Blur” (calibrated by TRACER)
 - Adding an additional midrange process blur (e.g. for HSQ) is possible (can be calibrated by TRACER)

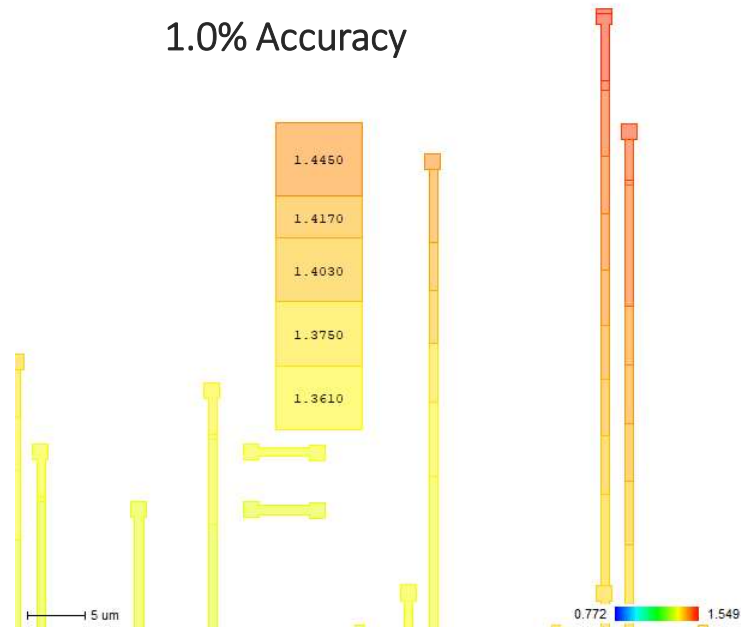


Dose Factor Accuracy

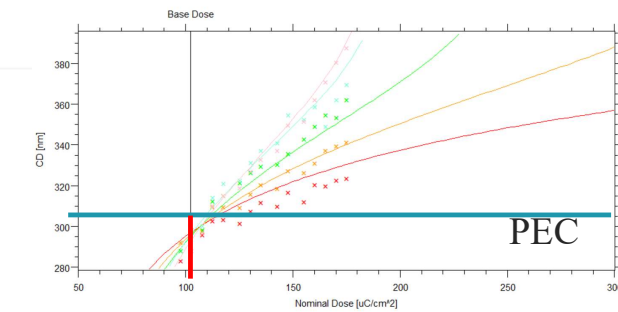
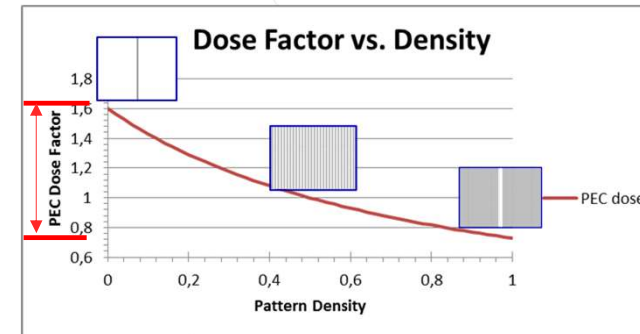
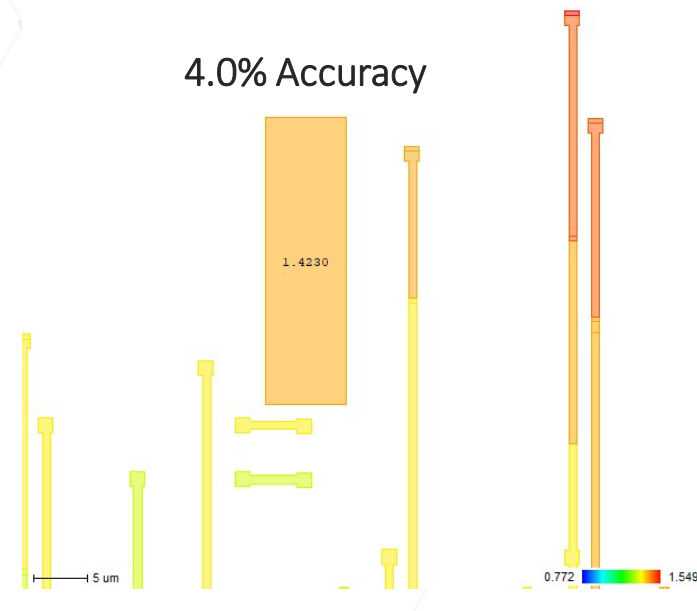
- The required dose range (PSF dependent) is split to “Dose Classes”
- Dose classes are automatically generated
 - Either via dose accuracy or manually by predefined dose classes
- Tradeoff between litho quality and shape count (write time)
 - Dose accuracy lower limited set by the system capability (typically 1%)



1.0% Accuracy

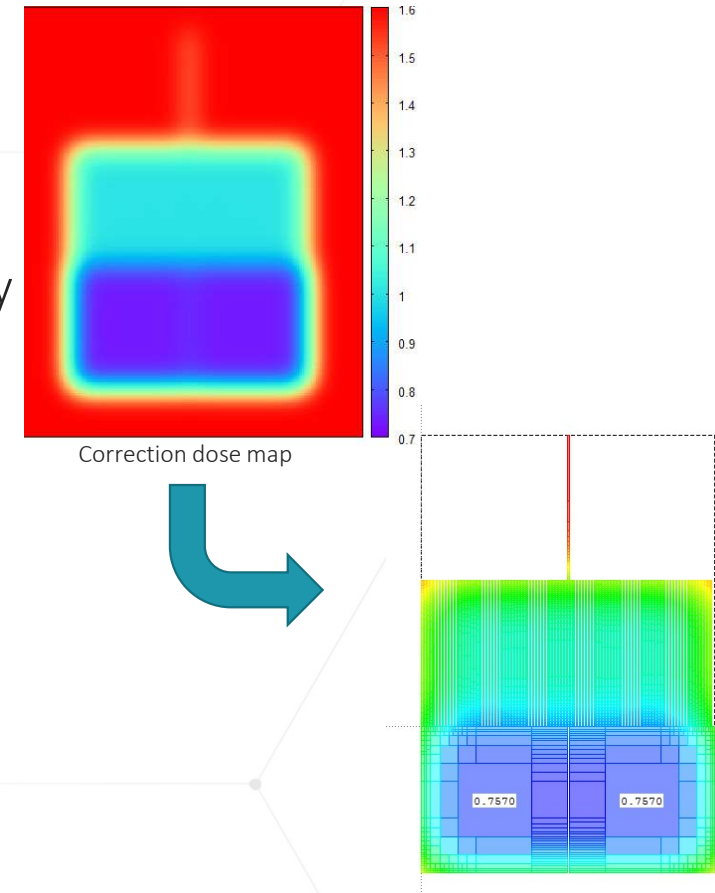
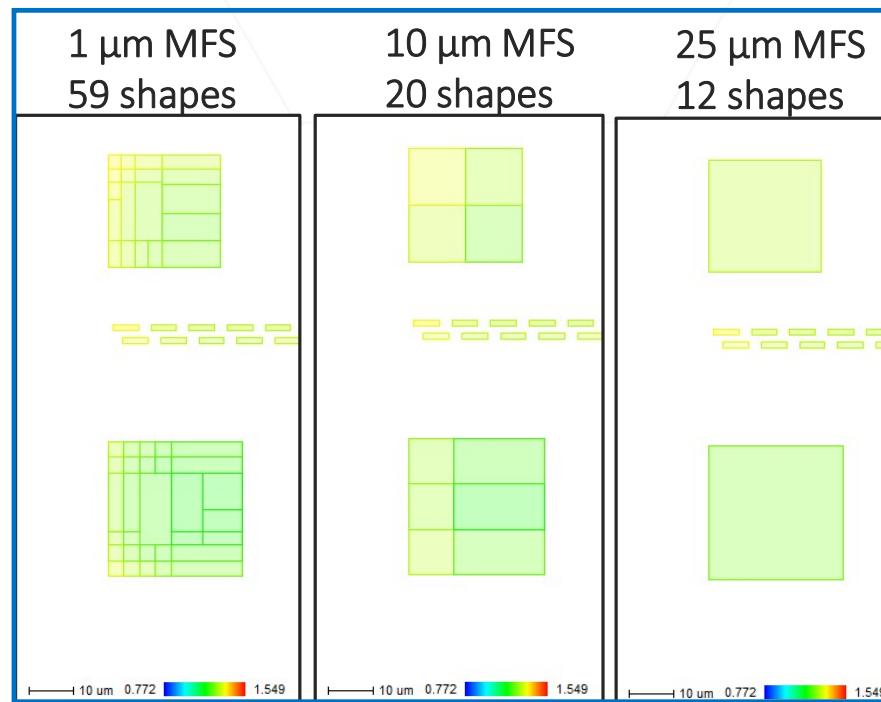
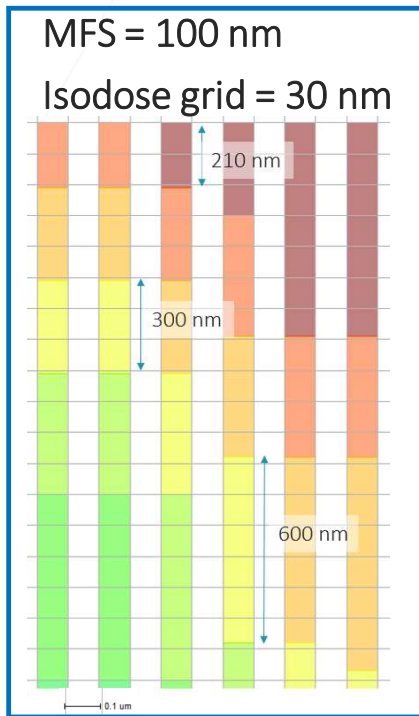


4.0% Accuracy



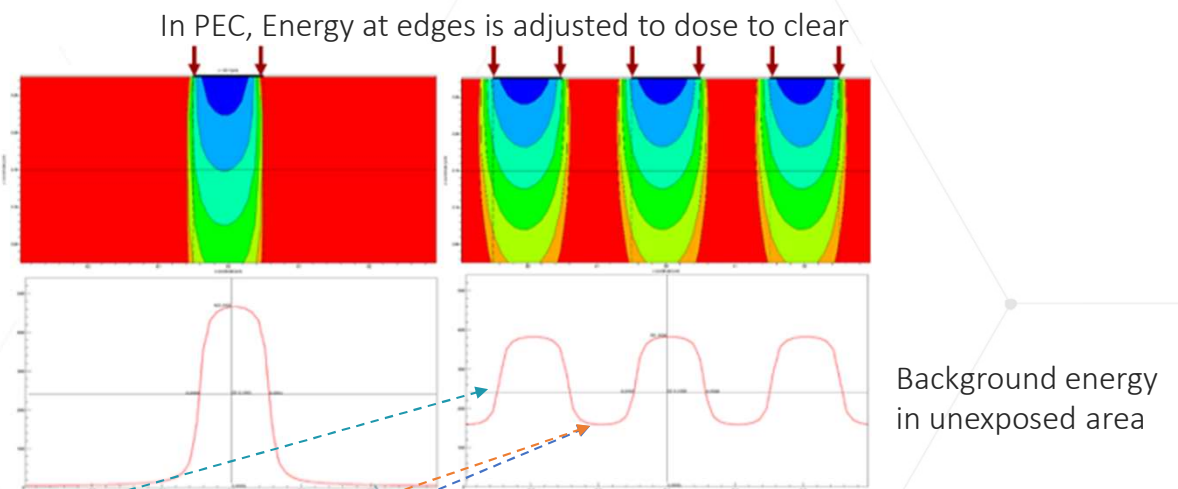
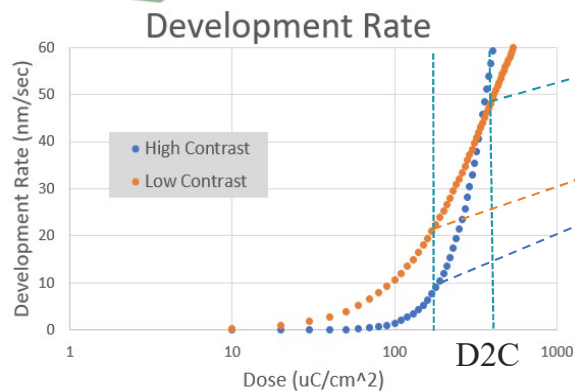
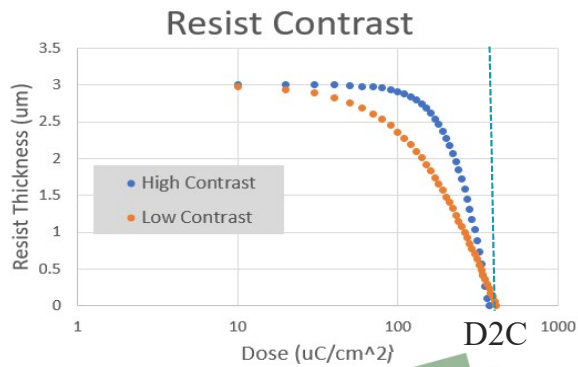
Physical PEC Fracturing

- The layout is locally fractured along the discretized dose map (iso-dose lines from dose classes)
 - “Iso-Dose” grid and Minimum Fracture Size (MFS) control location of cuts and number of shapes
 - Pre-fracturing allow to optimize number of shapes vs. accuracy



Origin of Lateral Development

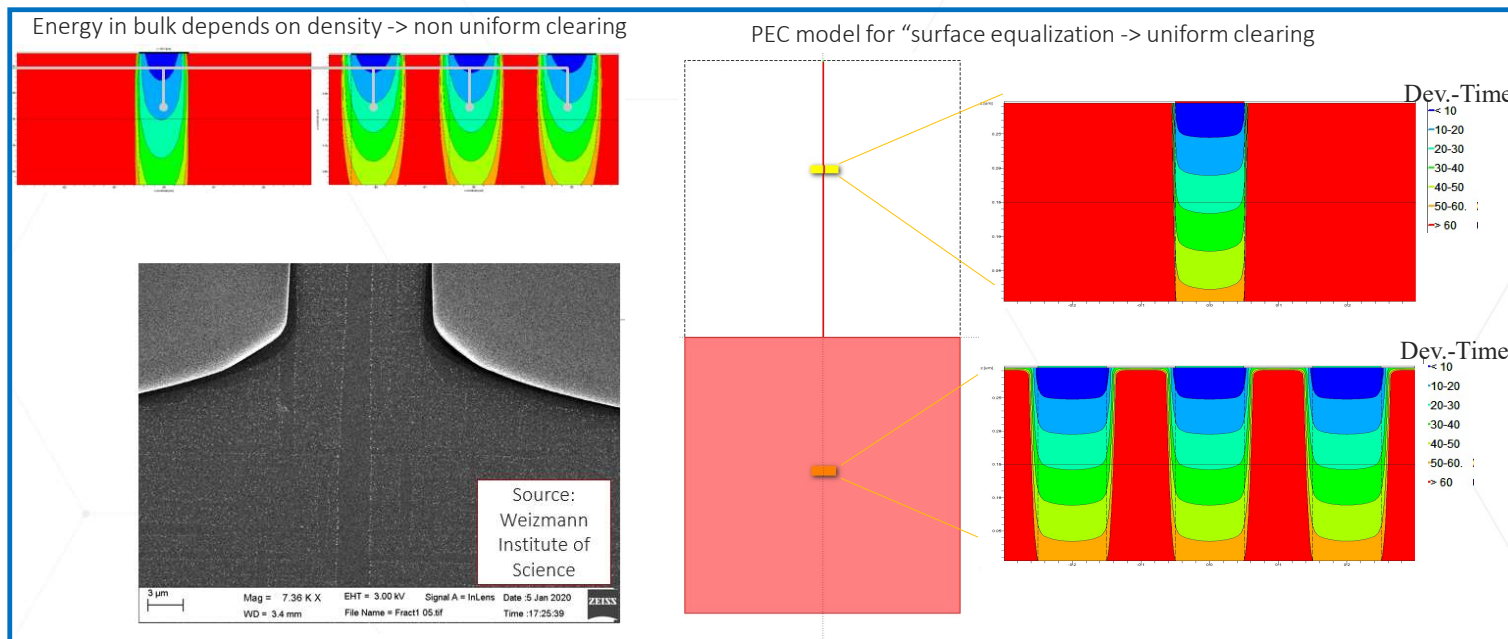
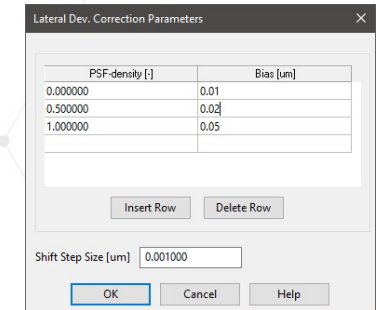
- Energy scatters beyond the pattern edges, leading to unintended partial exposure
- Practical resists have finite contrast; resist develops even for doses well below the dose to clear



- Together, these lead to development beyond exposed features, and features grow wider
- This is known as “Lateral Development”, and is an effect of the resist process

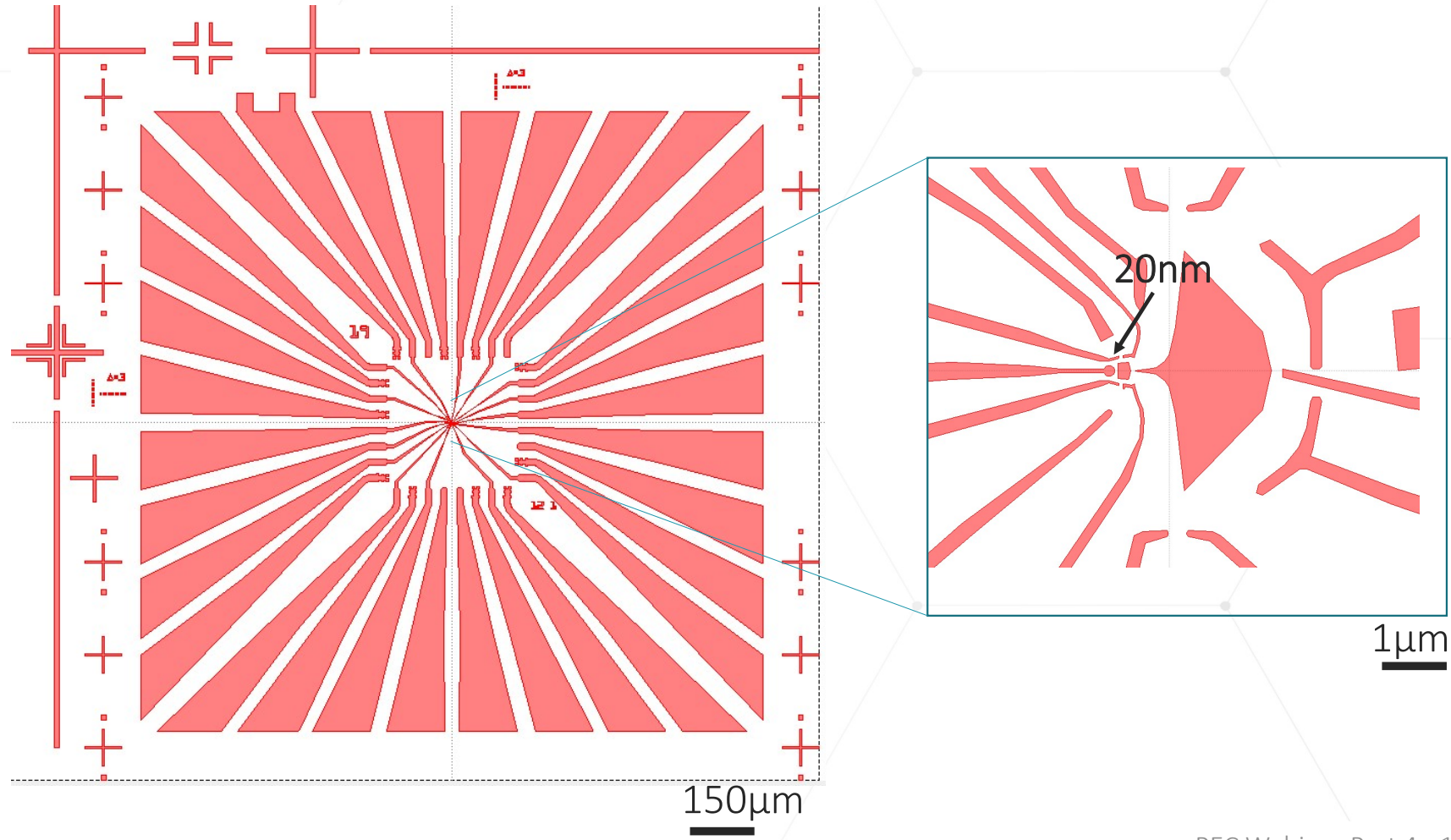
Advanced Process Correction Parameters

- Resist development effects require additional correction
- Lateral development is corrected by density-dependent bias
- Resist residues due to low energy in areas of high density (large pads) may require correction towards “uniform clearing”, a mix of OC/UC
- These are calibrated from experimental measurements using TRACER



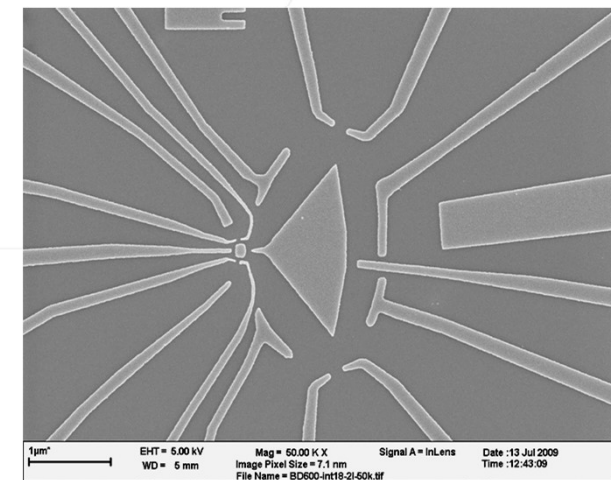
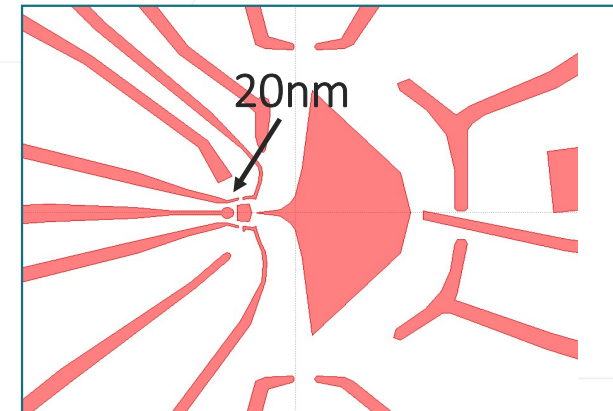
- Part 3 Summary: Dose PEC Parameter
- Process Effects and Major Parameter
 - From Design to Sample
 - Base Dose, Process Biases, Effective Blur and their Coupling
 - Calibration Strategy
- Calibration Procedure
- Advanced Model Parameter
- Summary
- Q&A

From Design To Sample



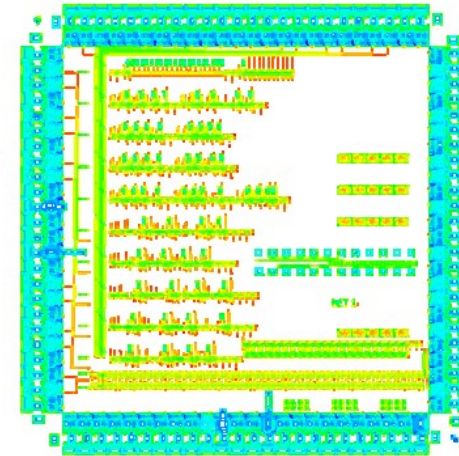
From Design To Sample

- What we see on SEM is the result of a complex process
 - Dataprep
 - Exposure
 - Writing strategy (fields, shape filling , ...)
 - Electron Scattering (spread of energy in 3D)
 - Resist development
 - Transfer of energy to dissolution rate
 - Resist development process (single layer, multi layer)
 - Post development process (baking, descum,...)
 - Pattern transfer
 - Lift off
 - Etching (wet, RIE)
 - Inspection
 - SEM imaging

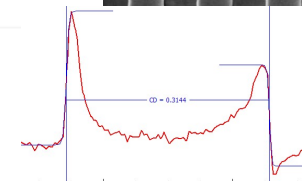
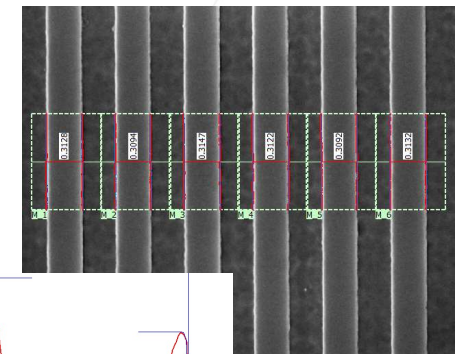


How do we Choose Exposure Parameters?

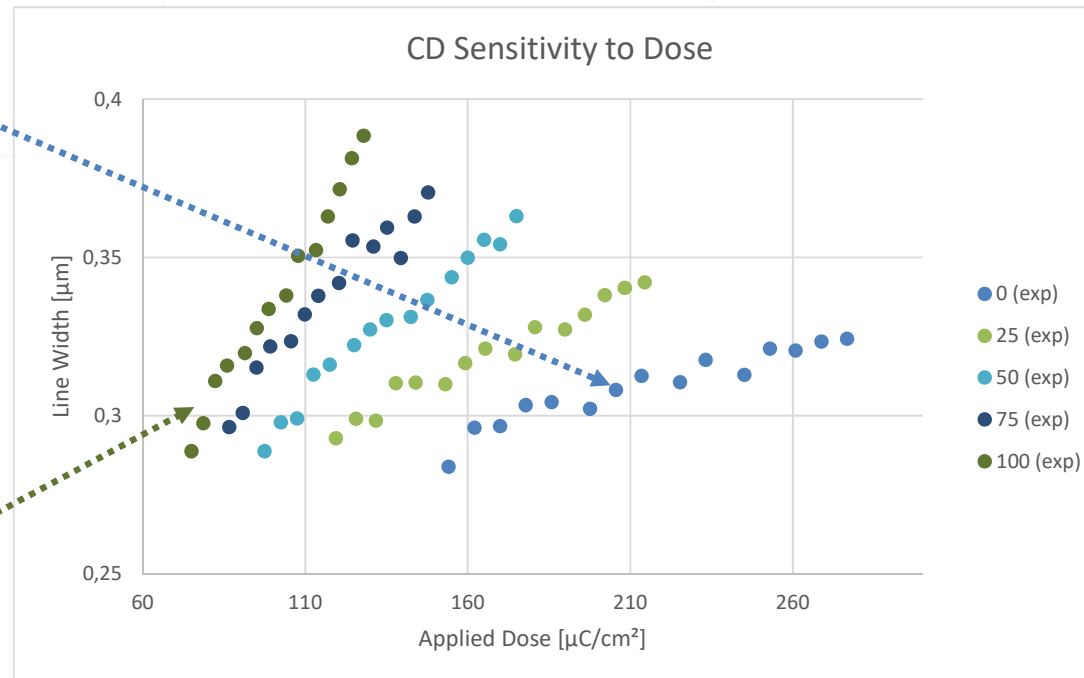
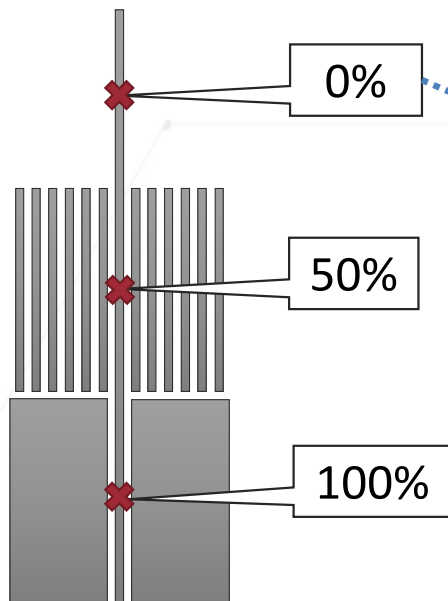
- Base Dose
 - Expose dose matrix
 - a „typical“ PEC'ed pattern
 - large 50% L/S pattern with dose matrix
 - Process (develop, pattern transfer)
 - Find „right dose“ with SEM measurement



- **Good as a starting point**
- **Does not include process effects across densities**
- **May not be the „best base dose“**



Typical Experimental Result



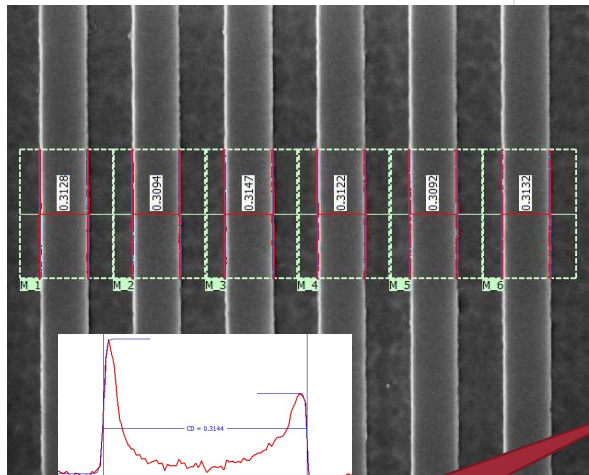
U.Penn:
200nm ZEP520A on Si

Accel. Voltage: 50 kV

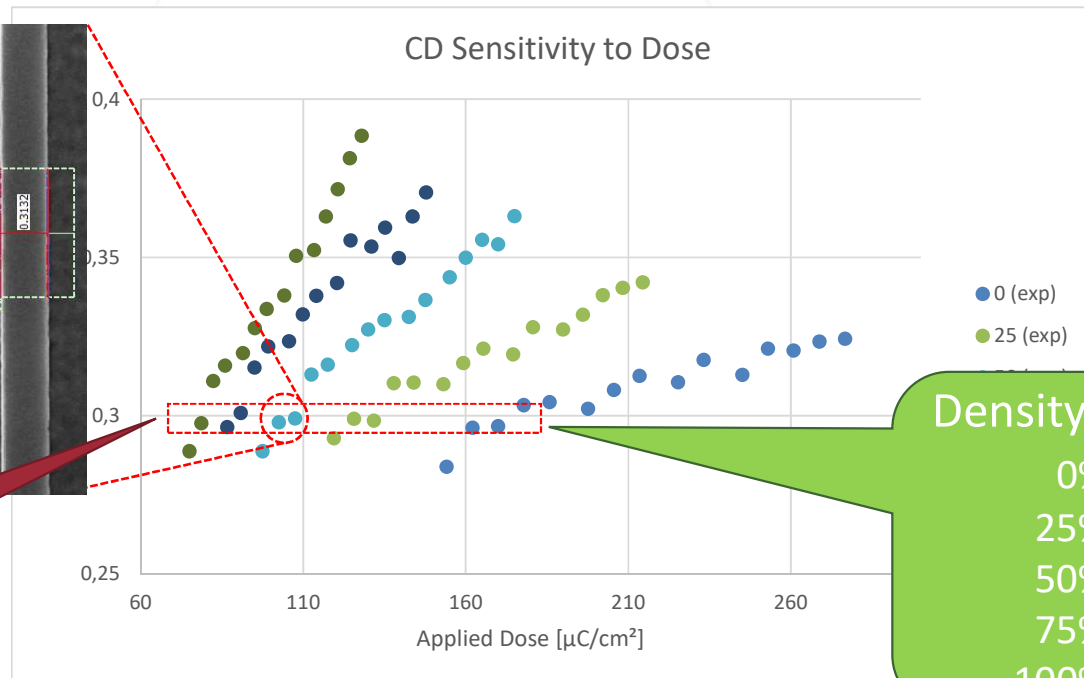
Target CD: 300nm

- For a given stack, CD_{measured} is a function of dose and pattern density
 - Iso- and dense lines require very different doses to get to the same CD
 - Iso lines show „best“ CD response with dose (big changes in dose \rightarrow CD variations)
- Base Dose and Bias is coupled

„Typical“ Process Optimization: Dose to Size



Target CD: 300 nm



U.Penn:
200nm ZEP520A on Si

Accel. Voltage: 50 kV

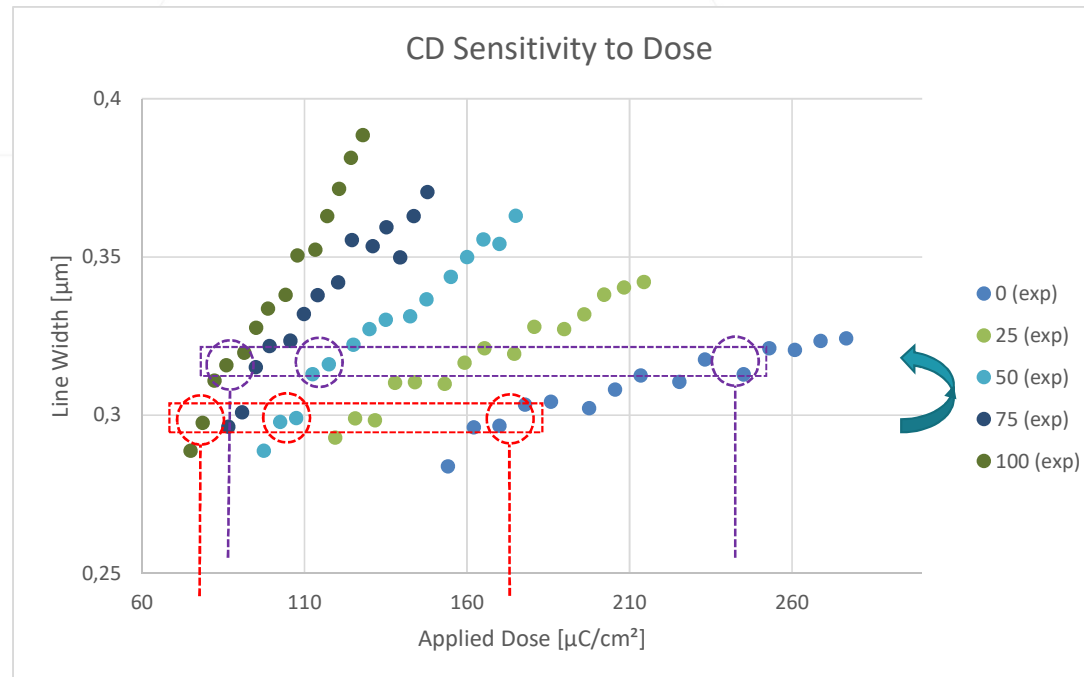
Target CD: 300nm

Density Dependent Doses

0%	170 $\mu\text{C}/\text{cm}^2$
25%	130 $\mu\text{C}/\text{cm}^2$
50%	107 $\mu\text{C}/\text{cm}^2$
75%	95 $\mu\text{C}/\text{cm}^2$
100%	80 $\mu\text{C}/\text{cm}^2$

- Typical base dose calibration picks „line width == space width“ for 1:1 L&S
 - Results in 107 $\mu\text{C}/\text{cm}^2$ Base Dose for this example
 - Results in Dose Range 80 (100%) .. 170 $\mu\text{C}/\text{cm}^2$ (0%)

Impact of „a“ Bias



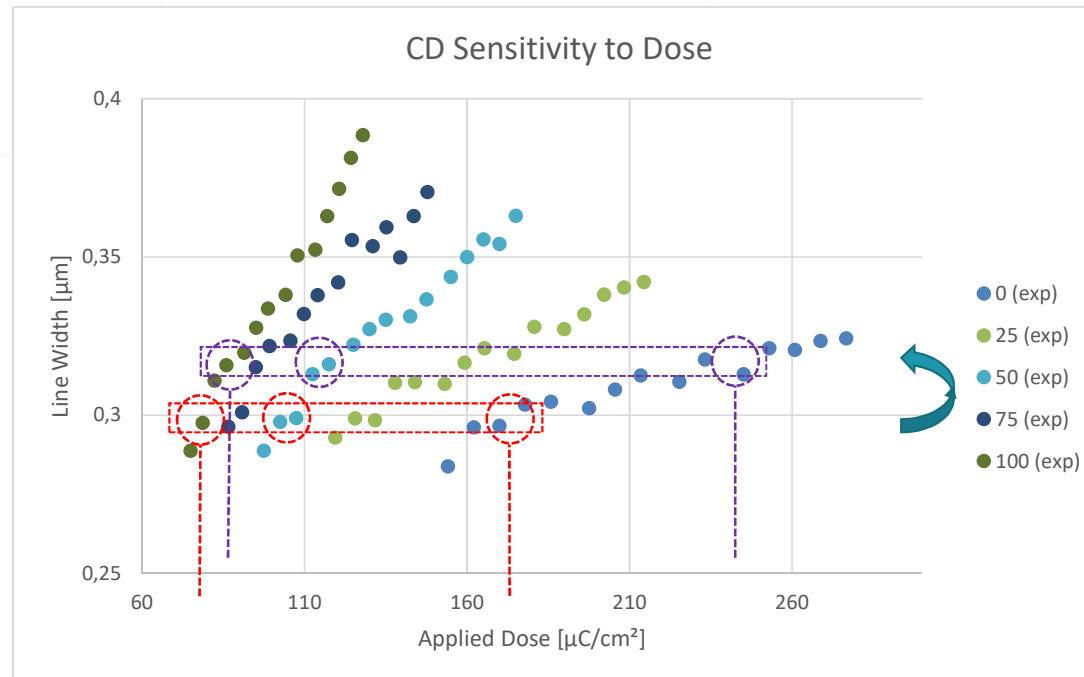
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200nm ZEP520A on Si

Accel. Voltage: 50 kV

Assume 20nm process bias
e.g. from development, etch, metrology

- Target CD (= Zero Bias) results in Dose Range 80 (100%) .. 170 $\mu\text{C}/\text{cm}^2$ (0%)
- 20nm Bias \Rightarrow Applied dose: 91 μC (100%) .. 233 $\mu\text{C}/\text{cm}^2$ (0%)

Impact of „a” Bias



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200nm ZEP520A on Si

Accel. Voltage: 50 kV

Assume 20nm process
bias
e.g. from development, etch,
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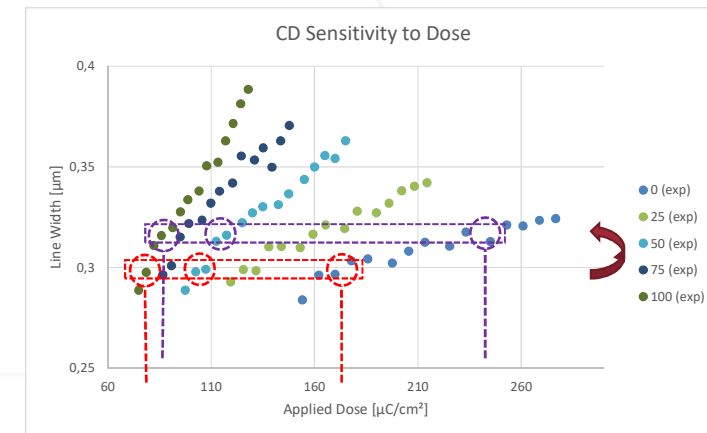
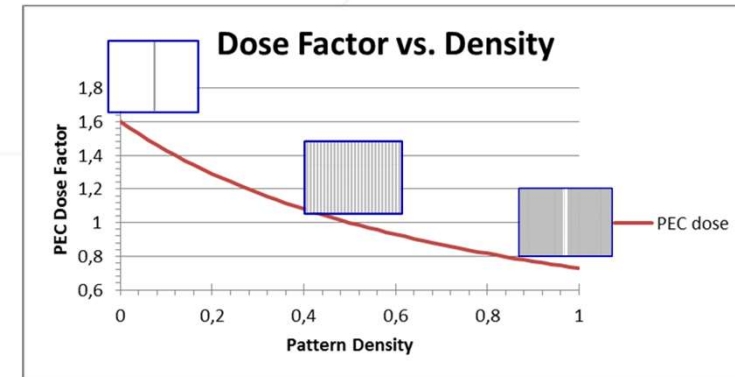
- Target CD (= Zero Bias) results in Dose Range 80 (100%) .. 170 µC /cm² (0%)
 - PEC would need to deliver a dose ratio $D_{iso} / D_{dense} = 170 / 80 = 2.12$
- 20nm Bias \Rightarrow Applied dose: 91µC (100%) .. 233µC /cm² (0%)
 - PEC would need to deliver a dose ratio $D_{iso} / D_{dense} = 233 / 91 = 2.56$

- PEC computes (density dependent) dose factors
 - The dose ratio D_{iso} / D_{dense} only depends on back-scattering (NOT on process point)
 - For Si at 50keV, $D_{iso} / D_{dense} = 2.4$

$$D_f = \frac{1}{1 + BE(2\rho - 1)} \quad \begin{array}{l} BE = 0.4 \\ \rho = 1 \text{ for dens / 0 isolated} \end{array}$$

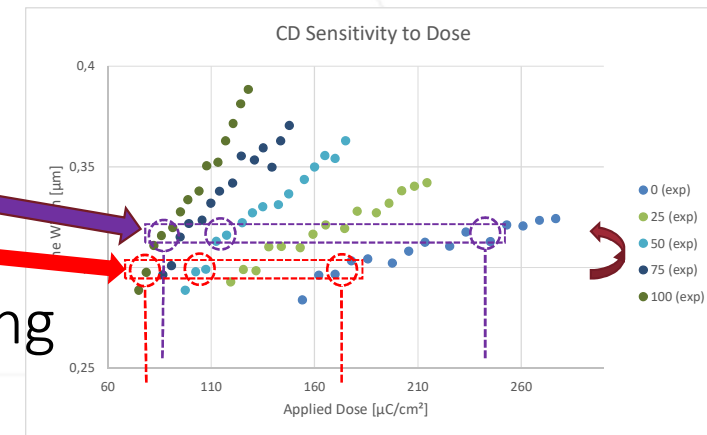
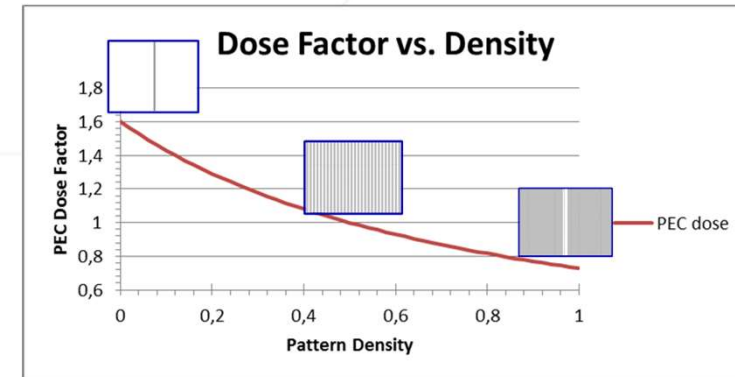
$$\frac{D_{iso}}{D_{dense}} = \frac{\frac{1}{1 + 0.412(2 * 0 - 1)}}{\frac{1}{1 + 0.412(2 * 1 - 1)}} = 2.4$$

Key Learning



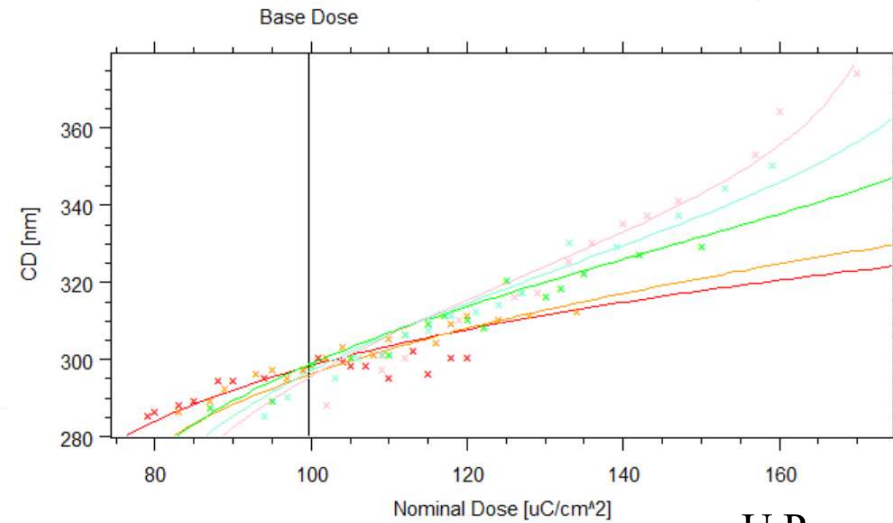
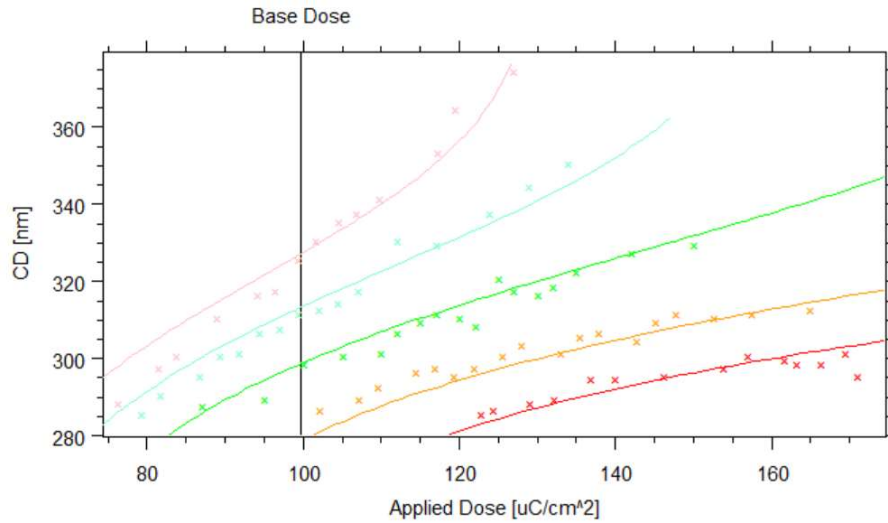
Key Learning

- PEC computes (density dependent) dose factors
 - The dose ratio D_{iso} / D_{dense} only depends on back-scattering (NOT on process point)
 - For Si at 50keV, $D_{iso} / D_{dense} = 2.4$
- However, process data shows varying dose ratios
 - For the sample data shown
 - $D_{iso} / D_{dense} = 2.56$ (20nm bias)
 - $D_{iso} / D_{dense} = 2.12$ (0 bias)
- Adjustment to proper dose range enable decoupling
 - Base Dose, Process Bias (global & density dependent), effective blur



Process point with smallest spot-size (blur) sensitivity: the iso-focal

Iso-Focal Calibration



U.Penn:
300nm ZEP520A on Si

Accel. Voltage: 50 kV

Applied Dose = Base Dose * D_f

Please Note: Both graphs show the same process!

Line Width vs. Applied Dose \Leftrightarrow **Line Width vs. Base Dose**

1. Chris Mack, Electron-beam lithography simulation for maskmaking, part IV, proceedings of Photomask and X-Ray Mask Technology VI, SPIE Vol. 3748, pp. 27-40
2. K.Keil et al, Determination of best focus and optimum dose for variable shaped e-beam systems by applying the isofocal dose method, Microelectronic Engineering 85 (2008) 778–781
3. U.Hofmann, N.Ünal, S.Sayan, G.Lopez, D.Mahalu, A novel method to find the best (iso-focal) process point in electron beam lithography, GenISys White Paper
4. G. Lopez et al, Isofocal Dose Based Proximity Effect Correction - Tolerance to the Effective Process Blur. Journal of Vacuum Science & Technology B 35, 06G505, 2017
5. Application Note in GenISys download area, Full Process Calibration using TRACER: Experimental Procedure

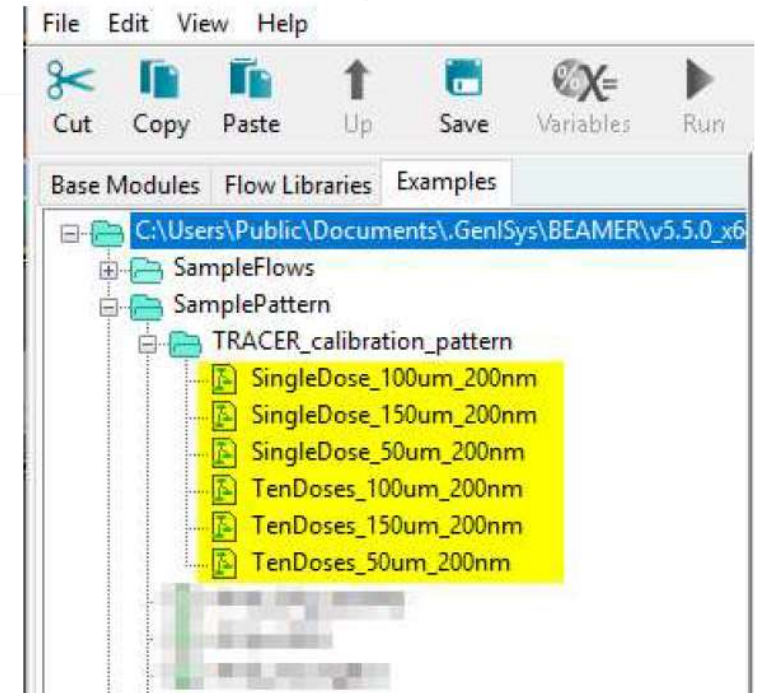
- Part 3 Summary: Dose PEC Parameter
- Process Effects and Major Parameter
- Calibration procedure
 - GaAs example
 - High contrast resist vs. low contrast resist
- Advanced Model Parameter
- Summary
- Q&A

- Calibration patterns are in BEAMER example folder
- Application Note in download area
- Help: support@genisys-gmbh.com



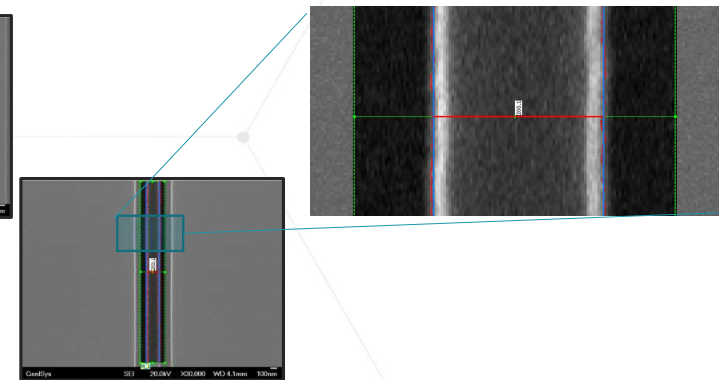
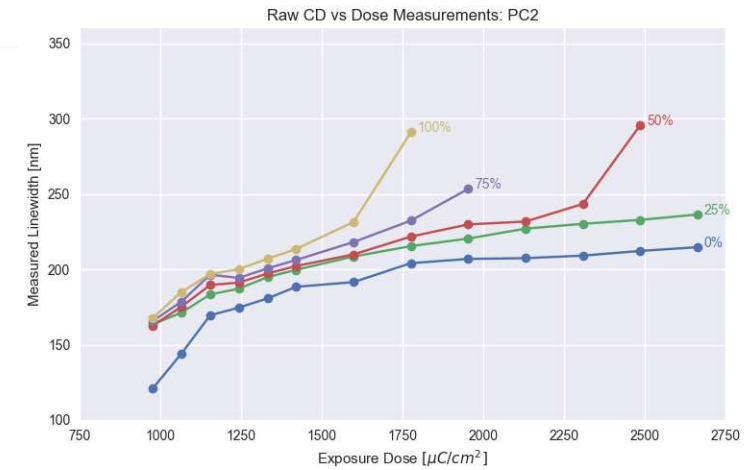
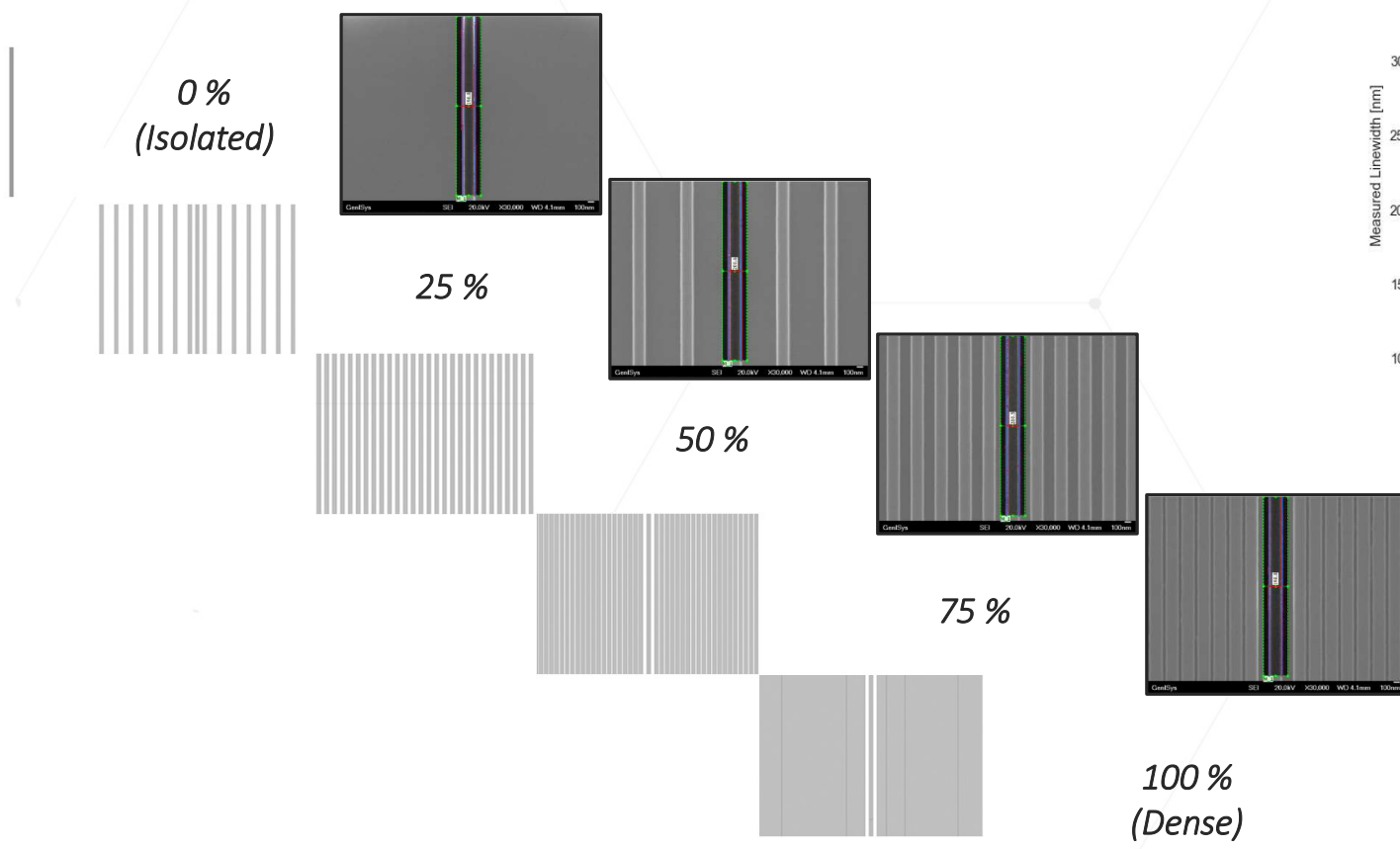
Full Process Calibration using TRACER: Experimental Procedure

An optimized e-beam lithography data preparation process must take into account process effects beyond just the electron energy distribution point spread function (PSF) as computed by TRACER. These process effects include density-dependent development rate changes, resist lateral development, and size bias due to process or metrology. It is possible to characterize and subsequently correct for these effects using a set of empirical measurements. This note describes the experimental procedure and data analysis necessary for such a Full Process Calibration.



Measuring Process Effects of Pattern Density

- Expose lines over a range of doses, with local pattern density varying from isolated to fully dense.



Fast and Consistent Measurements

- Consistent and reliable SEM measurements are critical for process calibration
 - Hand-drawn cursors are subjective, tedious, time-consuming, inconsistent
- ProSEM offers stable, consistent, fast CD measurements from saved SEM images
- Recipes, Batch Processing and Scripting enable automation

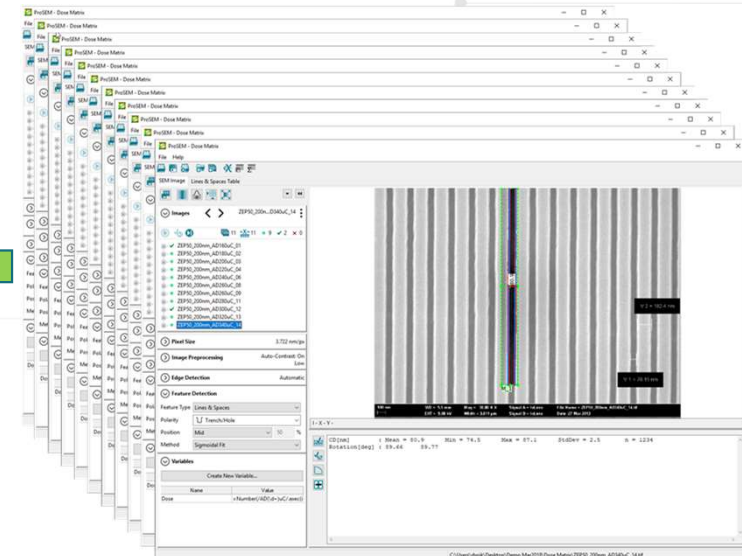


Automated SEM Image Analysis and Metrology Software



ProSEM - Dose Matrix

Image	Validation	Measurement ID	Dose	CD Mean[nm]	CD StdDev[nm]	CD Min[nm]	CD Max[nm]
ZEP90_200nm_AD160uC_01	Validated	M_1	160.0	54.6	2.2	48.0	61.7
ZEP90_200nm_AD180uC_02	Success	M_1	180.0	63.4	1.1	60.2	66.4
ZEP90_200nm_AD200uC_03	Success	M_1	200.0	67.8	1.1	64.3	70.9
ZEP90_200nm_AD220uC_04	Success	M_1	220.0	68.5	1.2	63.2	72.3
ZEP90_200nm_AD240uC_06	Success	M_1	240.0	70.2	1.2	67.1	73.6
ZEP90_200nm_AD260uC_08	Success	M_1	260.0	77.9	1.1	75.1	82.1
ZEP90_200nm_AD260uC_09	Success	M_1	260.0	77.9	1.1	75.1	82.1
ZEP90_200nm_AD280uC_11	Success	M_1	280.0	78.2	1.8	73.7	82.5
ZEP90_200nm_AD300uC_12	Validated	M_1	300.0	81.6	1.4	78.5	86.3
ZEP90_200nm_AD320uC_13	Success	M_1	320.0	79.8	1.0	76.9	82.5
ZEP90_200nm_AD340uC_14	Success	M_1	340.0	66.9	2.5	74.5	87.1



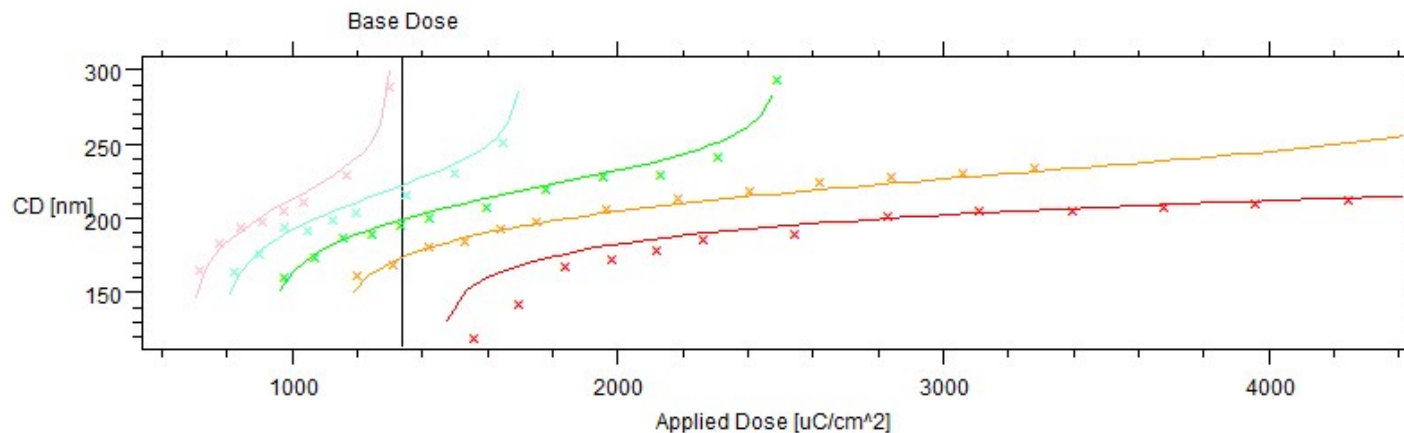
Fitting the Measured CD Data

- The data is fitted to determine additional correction terms needed to compensate for process effects

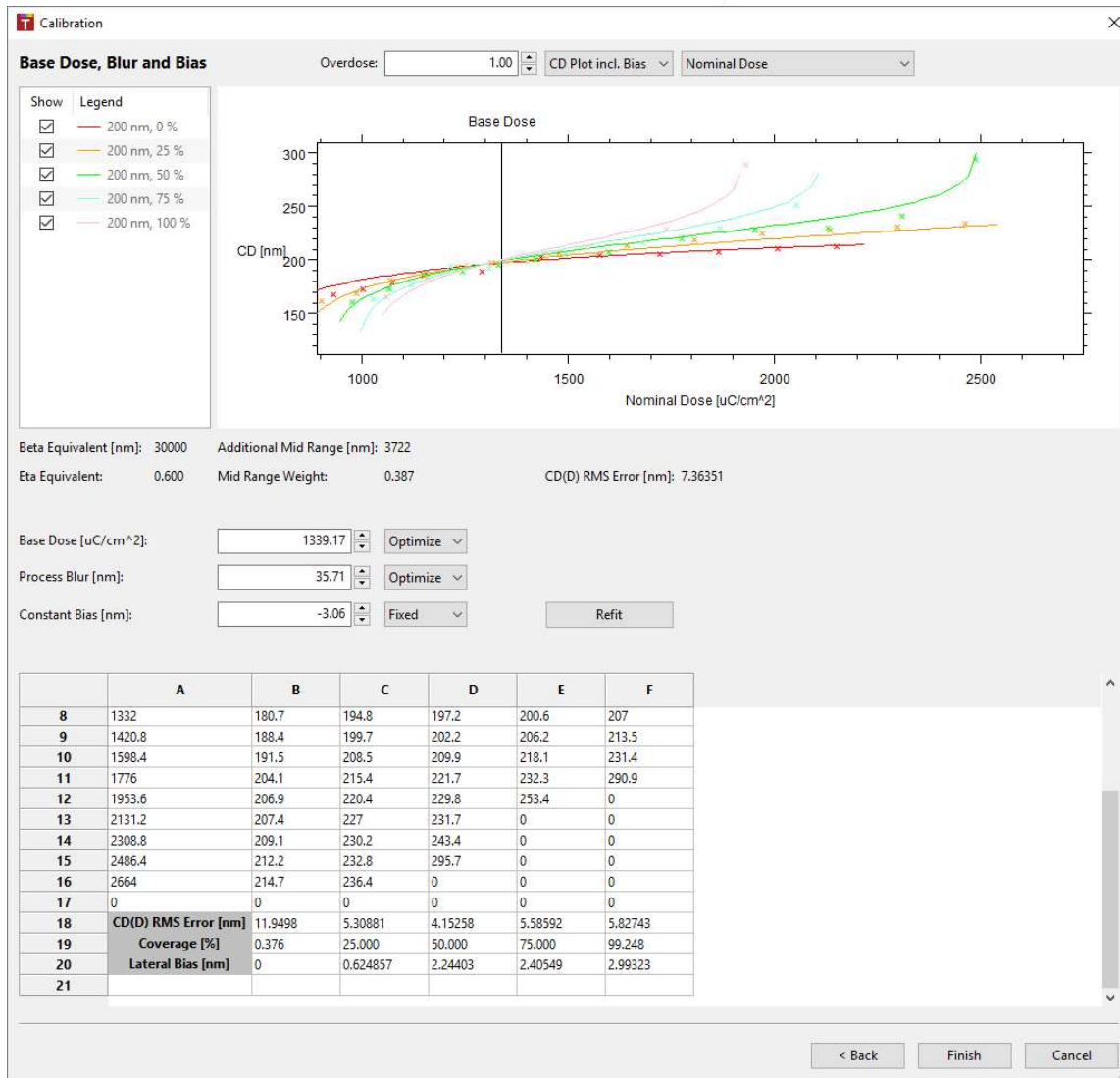
$$CD = CD_0 + Bias_{Lat.Dev} + \frac{ProcessBlur}{\sqrt{\ln(2)}} * Erf^{-1} \left[\left(\frac{D2C}{D} - BE - Dens_{long-range} * BE \right) * (1 - BE) \right]$$

Where the $Bias_{Lat.Dev}$ is determined using the development rates derived from the contrast curve and iterating the integral:

$$t_{lat} = \int_{x_0}^{x_1} \frac{dx}{r_{lat}(x)}$$

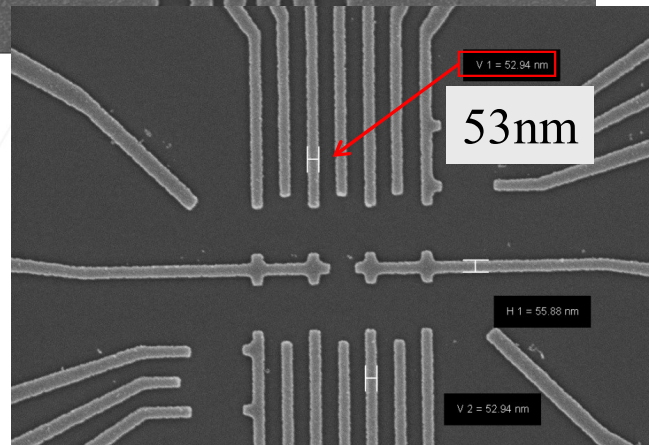
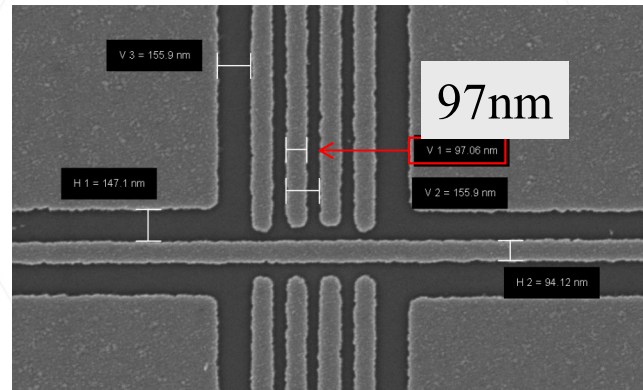
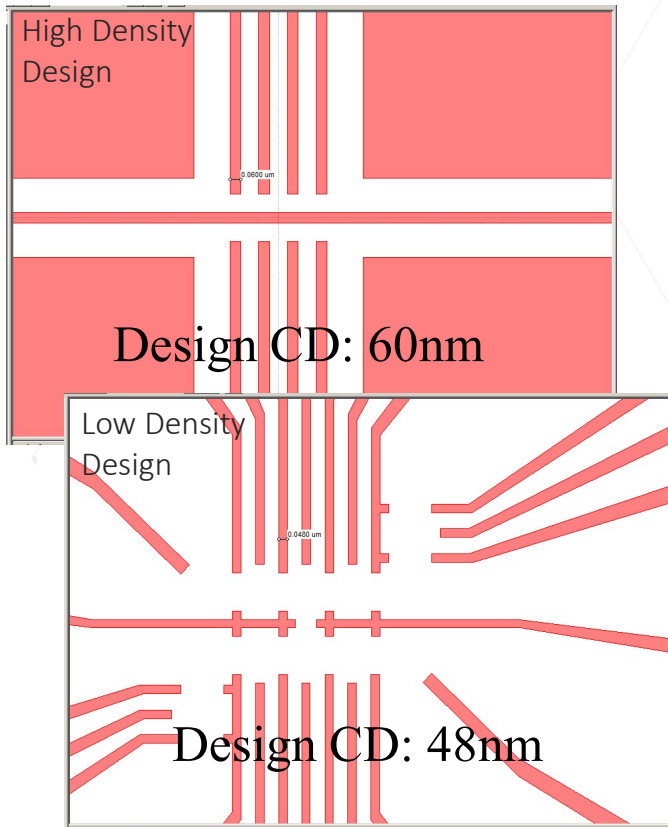


Fit Results



- The fitting procedure results in an “Extended Point Spread Function”, adding terms to the scattering PSF for:
 - Optimal Base Exposure Dose
 - Process Blur
 - Overall Process Bias
 - Density-dependent Bias to compensate for lateral development
 - Midrange Gaussian term for additional process effects, such as diffusion
 - OC/UC Mix Factor

Quantum Device w/o Process Calibration

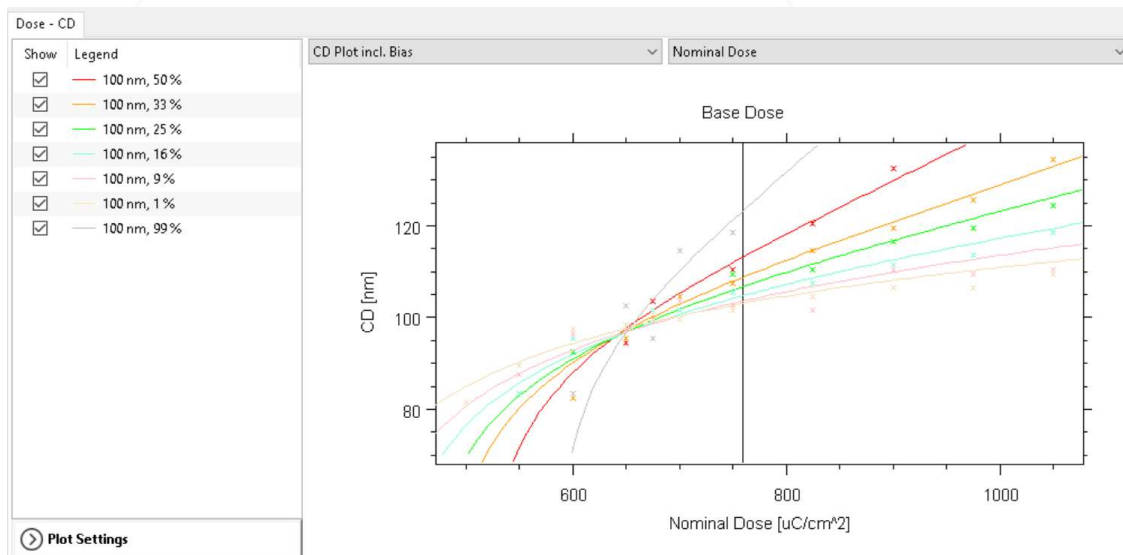


Conventional PEC Only

- The 48 nm line almost on target – 60nm line far away from target

• Calibration of Process Data resulted in

- Base Dose = 795 $\mu\text{C}/\text{cm}^2$
- Process Blur = 26nm
- Bias_{0%} = 4nm; Bias_{25%} = 9nm;
Bias_{50%} = 18nm; Bias_{99%} = 32nm

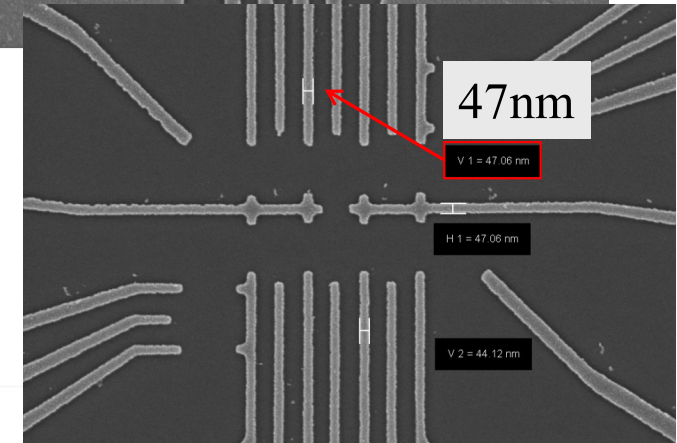
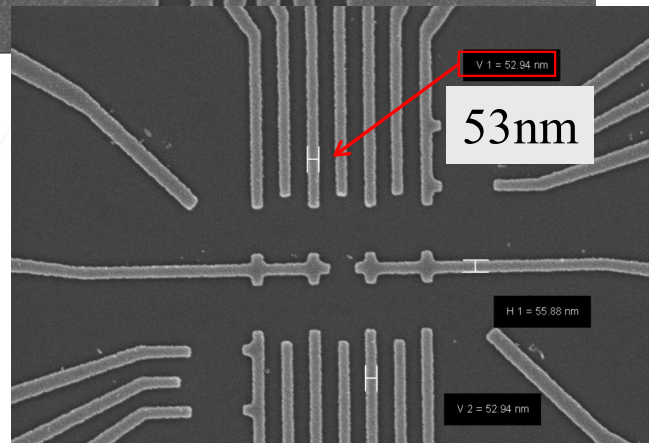
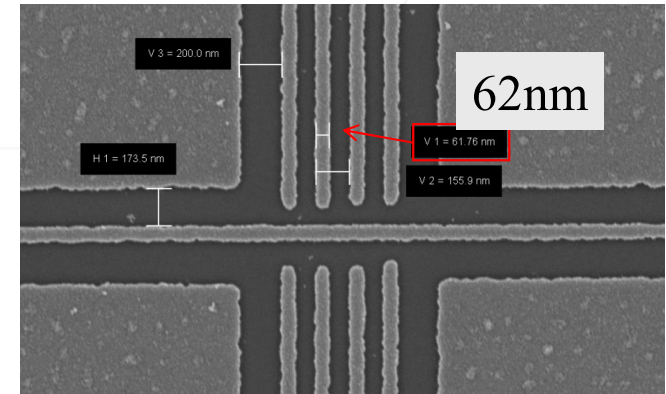
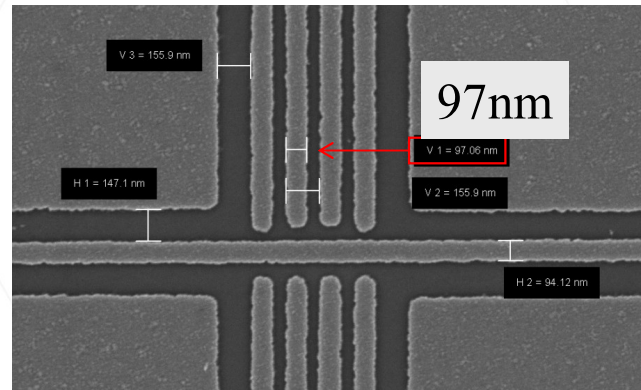
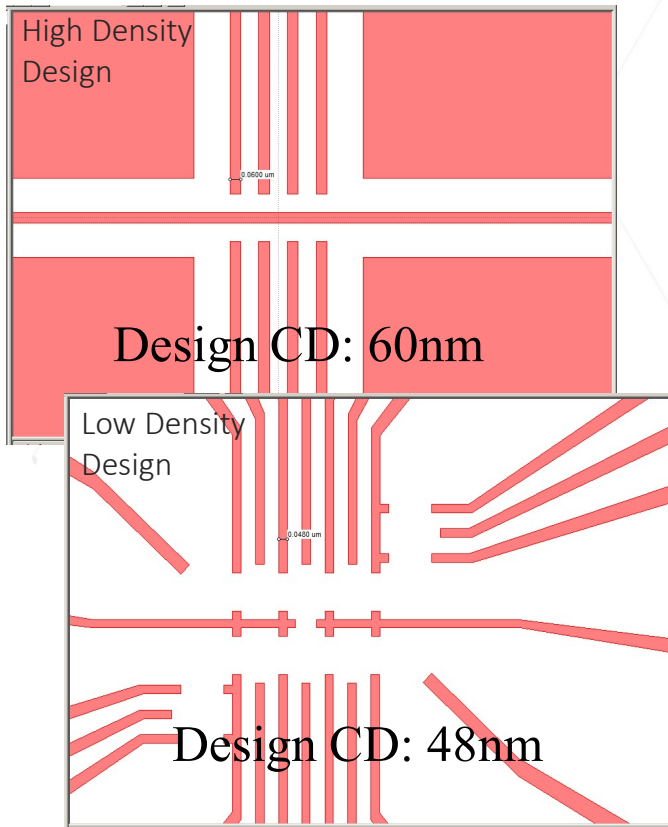


Calibration for 100keV on GaAs

12°C (tool set point) development 2 minutes								
Dose factor	Pattern Ratio	1:1	1:2	1:3	1:5	1:10	1:1000	100:1
1050 $\mu\text{C}/\text{cm}^2$		--	135.6	125.9	119.1	111.3	110.8	--
975 $\mu\text{C}/\text{cm}^2$		--	126.2	120.6	114	110.8	107.4	--
900 $\mu\text{C}/\text{cm}^2$		132.9	120.9	117.6	112.1	111.8	107.9	--
825 $\mu\text{C}/\text{cm}^2$		121.6	115.5	111.5	108.5	102.5	105	--
795 $\mu\text{C}/\text{cm}^2$		118	112	110	106	105	104	132
750 $\mu\text{C}/\text{cm}^2$		111.5	108.2	110	105.6	102.9	102.5	118.7
700 $\mu\text{C}/\text{cm}^2$		104.7	104.7	104.1	102	103.7	100.4	115.1
675 $\mu\text{C}/\text{cm}^2$		104.1	100.9	100.3	102.2	100	101	95.6
650 $\mu\text{C}/\text{cm}^2$		95	95.7	99.1	103.4	97.8	98.9	103.7
600 $\mu\text{C}/\text{cm}^2$		81.6	89.7	93	95.6	94.1	98.1	--
550 $\mu\text{C}/\text{cm}^2$					83.8	88.2	90.4	
500 $\mu\text{C}/\text{cm}^2$							82.5	

Process:
200nm PMMA on GaAs
Exposure @ 100 kV
Development: 2 minutes at 12°C (tool set point)

Quantum Device Result



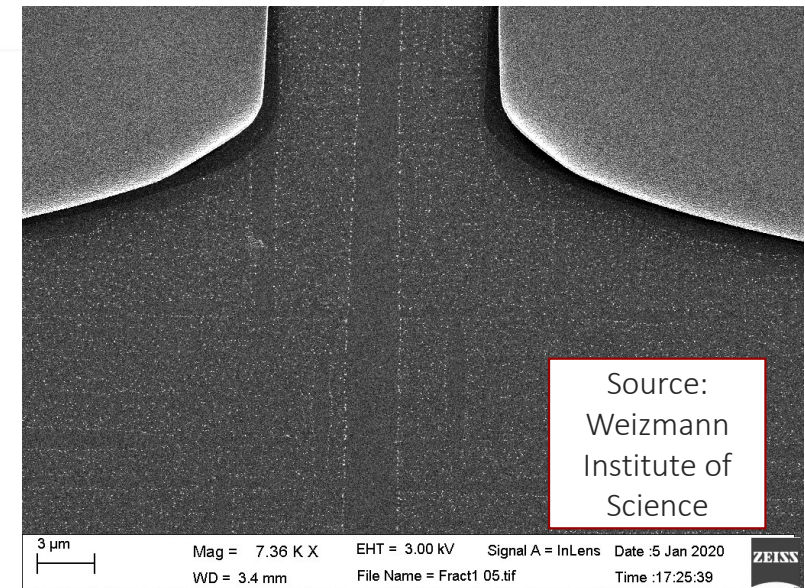
Conventional PEC Only

PEC with Process Calibration

- Excellent Results on different CDs at different densities

- In some cases, resist residues are left in large exposed areas
- This is especially found with:
 - Low contrast resists
 - On Higher Z materials, such as III-V semiconductor substrates

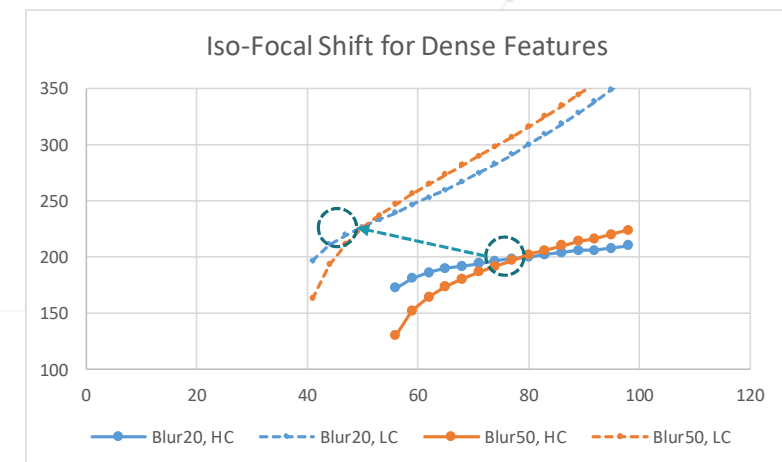
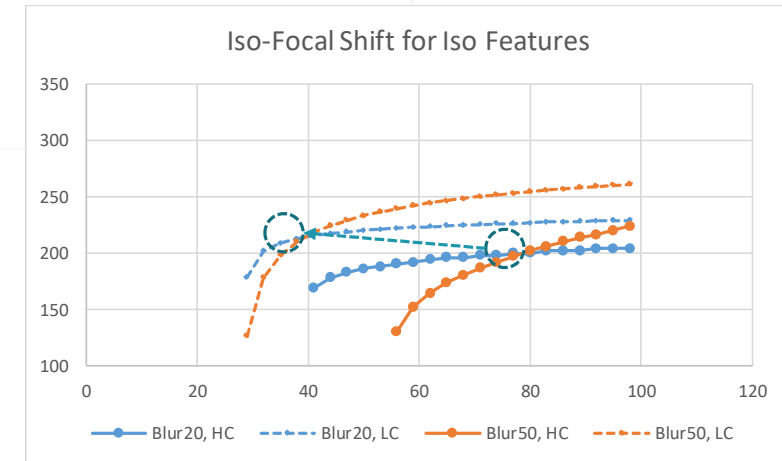
Resist Residues



- Part 3 Summary: Dose PEC Parameter
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 - HSQ peculiarities
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Lateral Development \Rightarrow Iso-Focal Shift

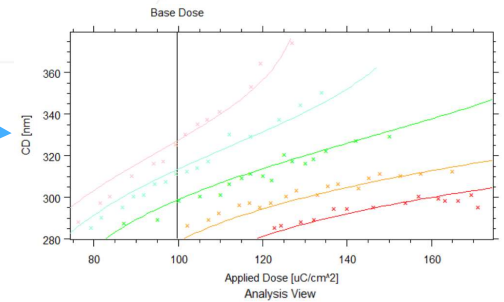
- Low-contrast resists develop also at lower doses
 - Iso: development into the blur (e.g. spot size)
 - Si example: 80 μ C -> 40 μ C, 12nm Bias
 - Dense: development into the blur and backscatter
 - Si example: 80 μ C -> 50 μ C, 25nm Bias
- Net effect: the process iso-focal shifts to lower doses
 - Stronger with more back-scattering (III-V materials)
 - Stronger with lower contrast resists (e.g. PMMA)
 - Stronger for thicker resist
- Key Learnings
 - For III-V on GaAs, this can shift the iso-focal below D2C
 - Since the amount of shift is density dependent, it will change the required PEC dose range



Blur-Dose Matrix

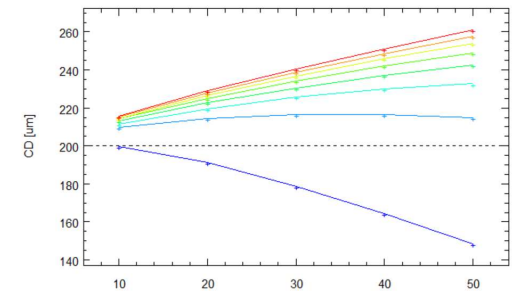
- Different ways to plot CD / Dose / Density / Blur Dependency

- CD as function of Dose, with Density Iso-Lines



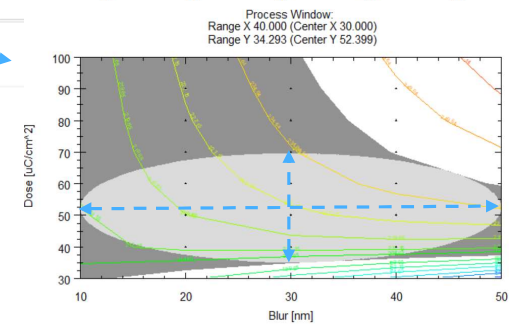
- CD as function of Blur, with Dose Iso-Lines

- Individual plots for the different densities
 - Already gives an indication of „iso-focal“ dose (horizontal trend)

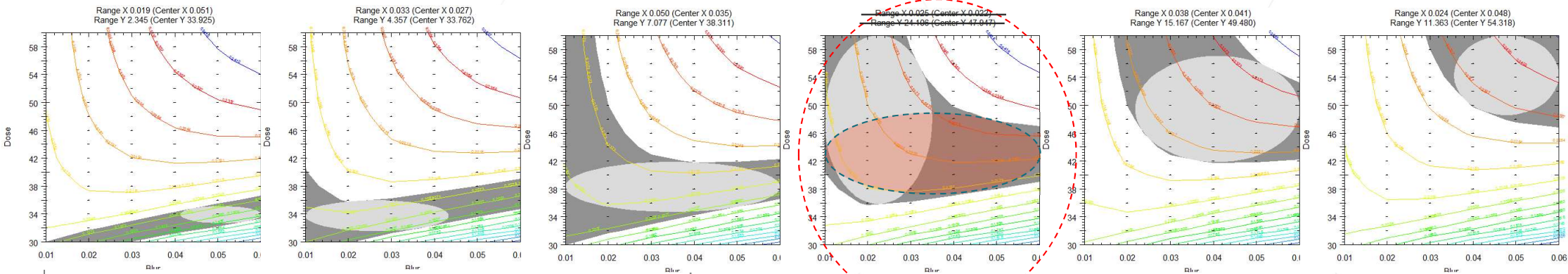


- Dose as a function of Blur, with CD iso-Lines

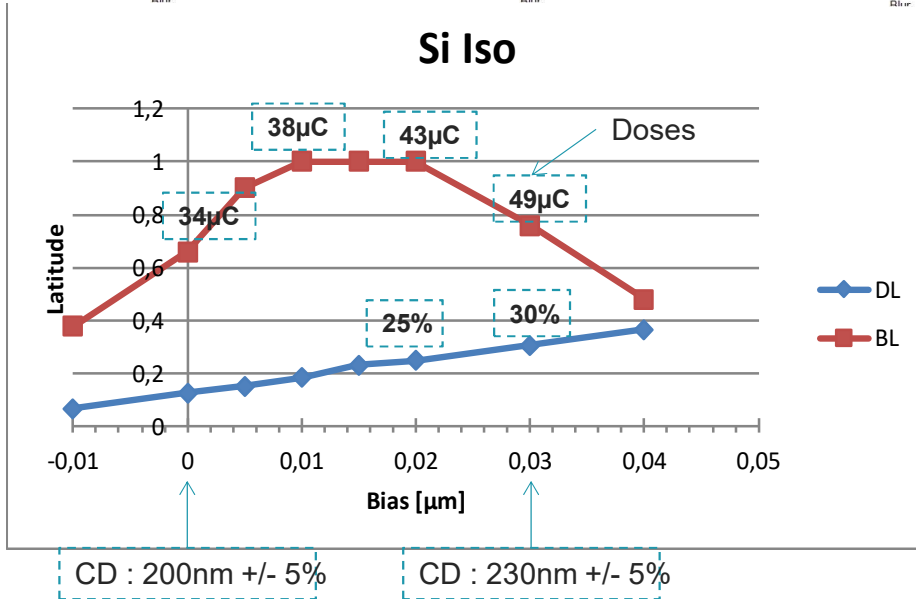
- $\pm 5\%$ CD tolerance from target provides CD limits (gray area)
 - Fit elliptical Process Window into CD limits
 - Horizontal axis is „blur latitude“
 - Vertical axis is „dose latitude“



Concept Study



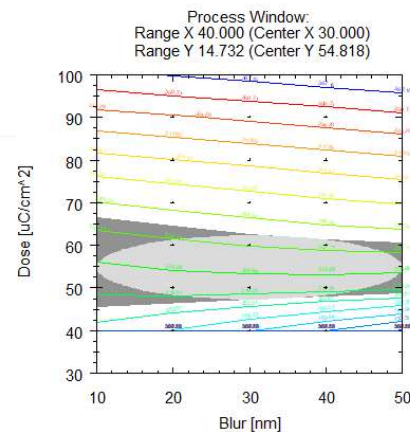
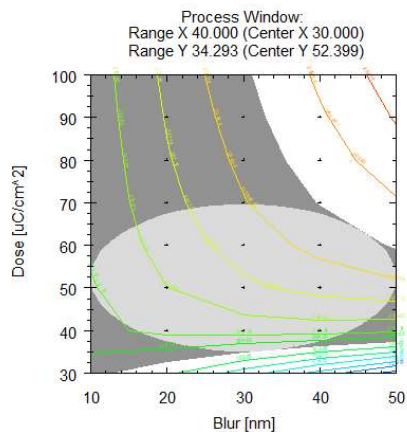
Si Iso



- Iso-Line, Si, Low-contrast Resist
- „Best“ Process Point @ $\sim 20\text{nm}$ Bias
 - „Largest“ Process Window
 - Smallest Bias Value
 - Base Dose = $40\mu\text{C}$

Process Iso-Focal

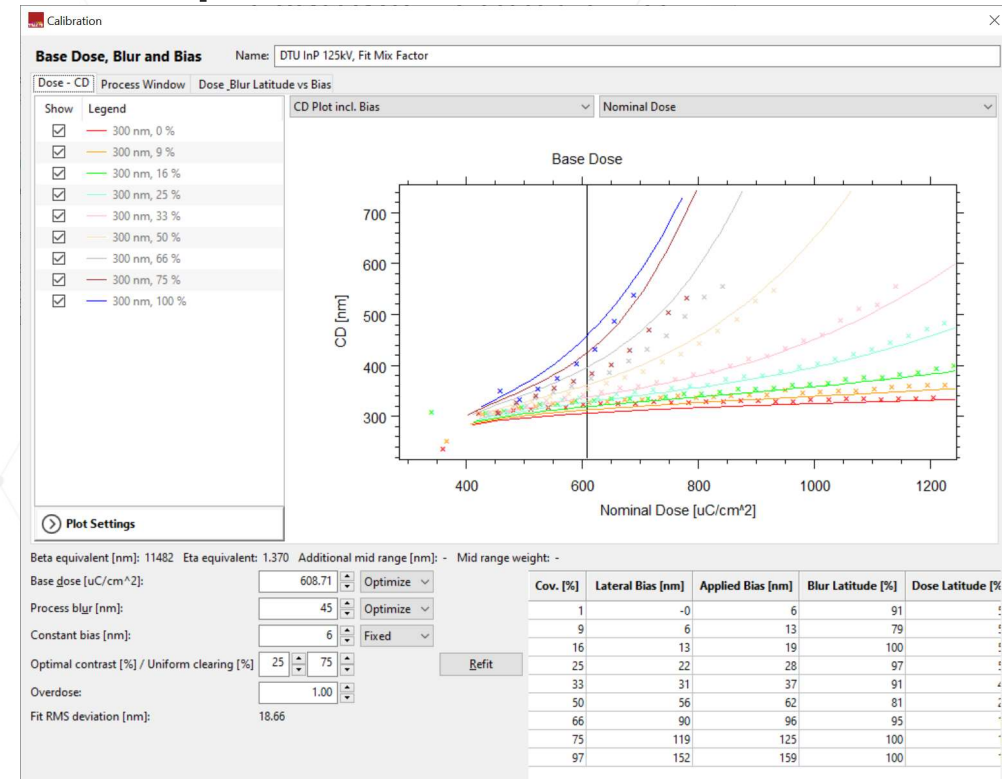
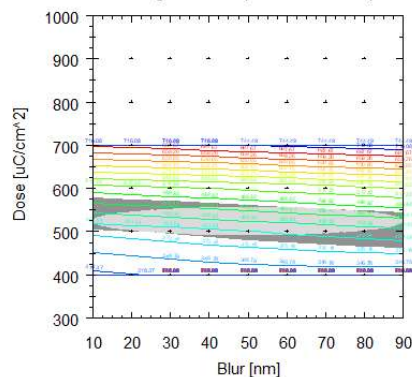
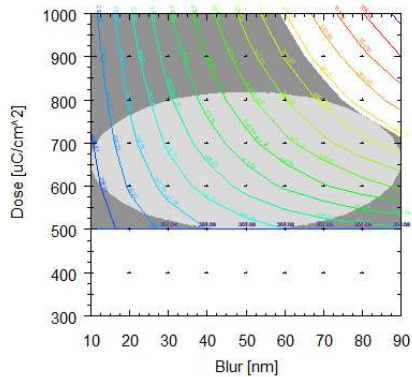
- For each density, search a large enough „blur latitude“
 - This guarantees good CD control also at the corners of the field
 - In the high-contrast case, this is equivalent to the optical iso-focal
 - TRACER searches for a large enough blur latitude (up to $2.5 * \text{ProcessBlur}$)
 - Under constraints ($\text{BaseDose} * \text{PEC_Dose}_{100\%} > \text{D2C}$, small Bias values)



- Please Note: blur cross-over (= iso-focal) and density cross-over are different points
 - Therefore, a density dependent Bias becomes essential

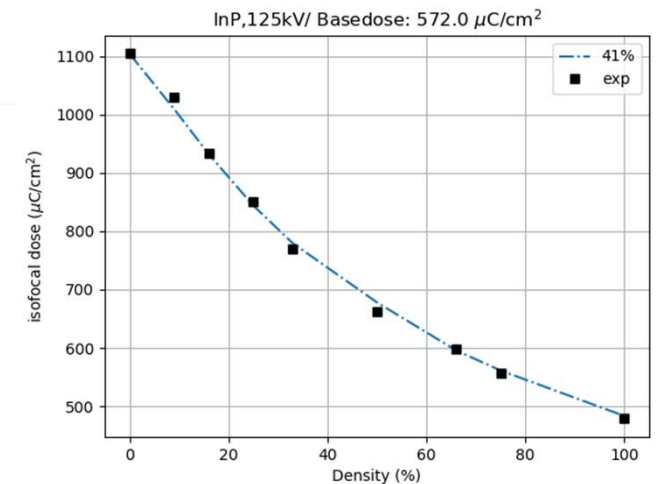
Low Contrast InP Example, Process Iso-Focal

- Customer Case
 - InP, 125kV, PMMA ($\gamma=3$)
 - D2C known as $480\mu\text{C}/\text{cm}^2$
- Process Iso-focal at $609\mu\text{C}/\text{cm}^2$
 - Mix-Factor at 25/75
 - Above D2C ($609 \cdot 0.88 = 538\mu\text{C}/\text{cm}^2$)



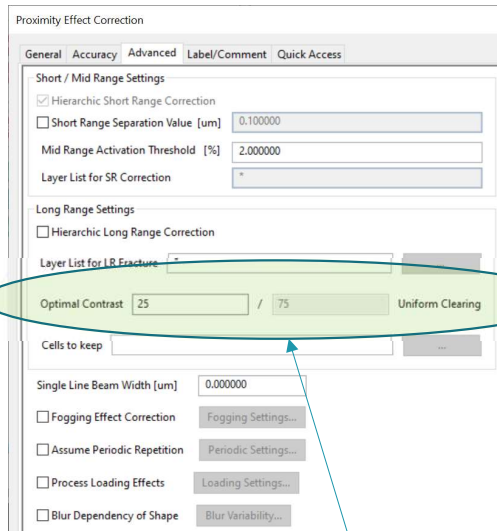
Low Contrast InP Validation

- Customer measured $572\mu\text{C}/\text{cm}^2$
 - Mix-factor of 41/59
- Pretty close to model prediction at $609\mu\text{C}/\text{cm}^2$
 - Mix-Factor of 25/75
- Not bad for just one dataset with one spot-size
 - Measuring process iso-focal requires at least two data sets with different spot sizes
 - TRACER criteria optimizes for „large enough“ blur latitude

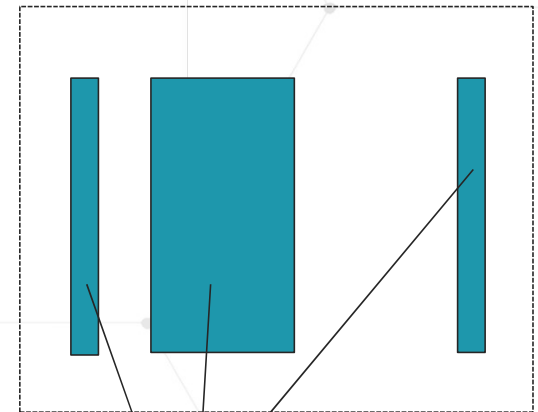
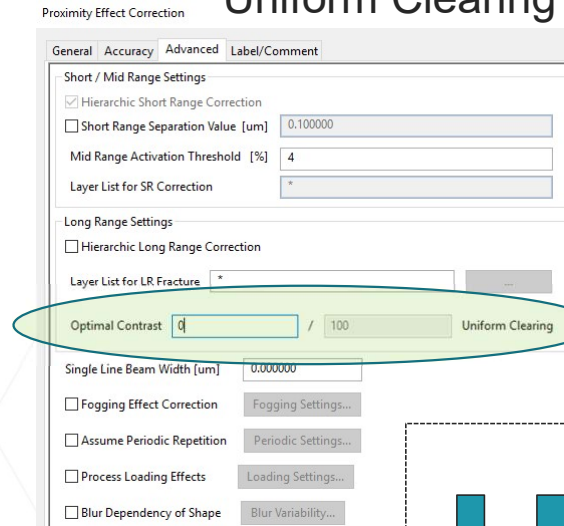


PEC – Mix Factor

Uniform Clearing = Surface Equalization



Automatically transferred via extended PSF

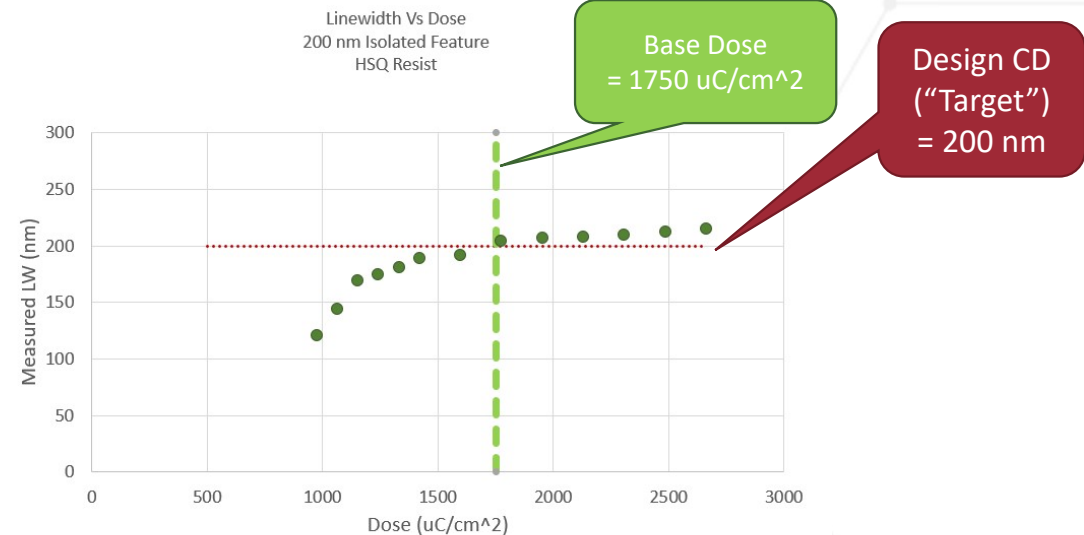
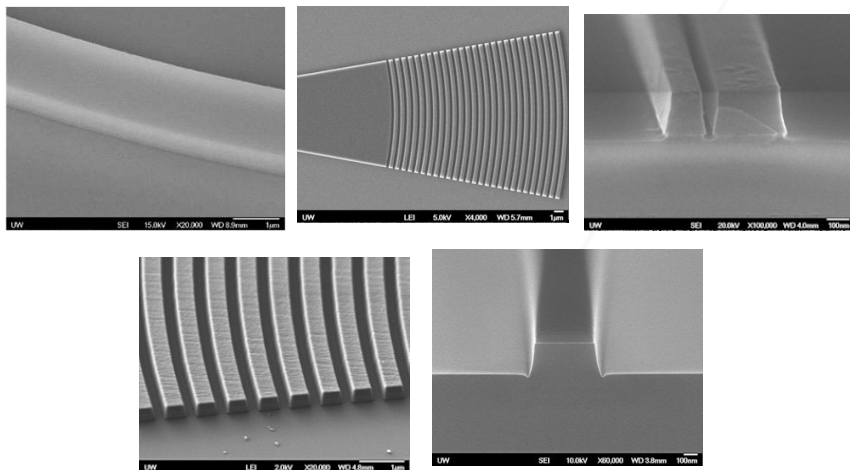


$$E = 1$$

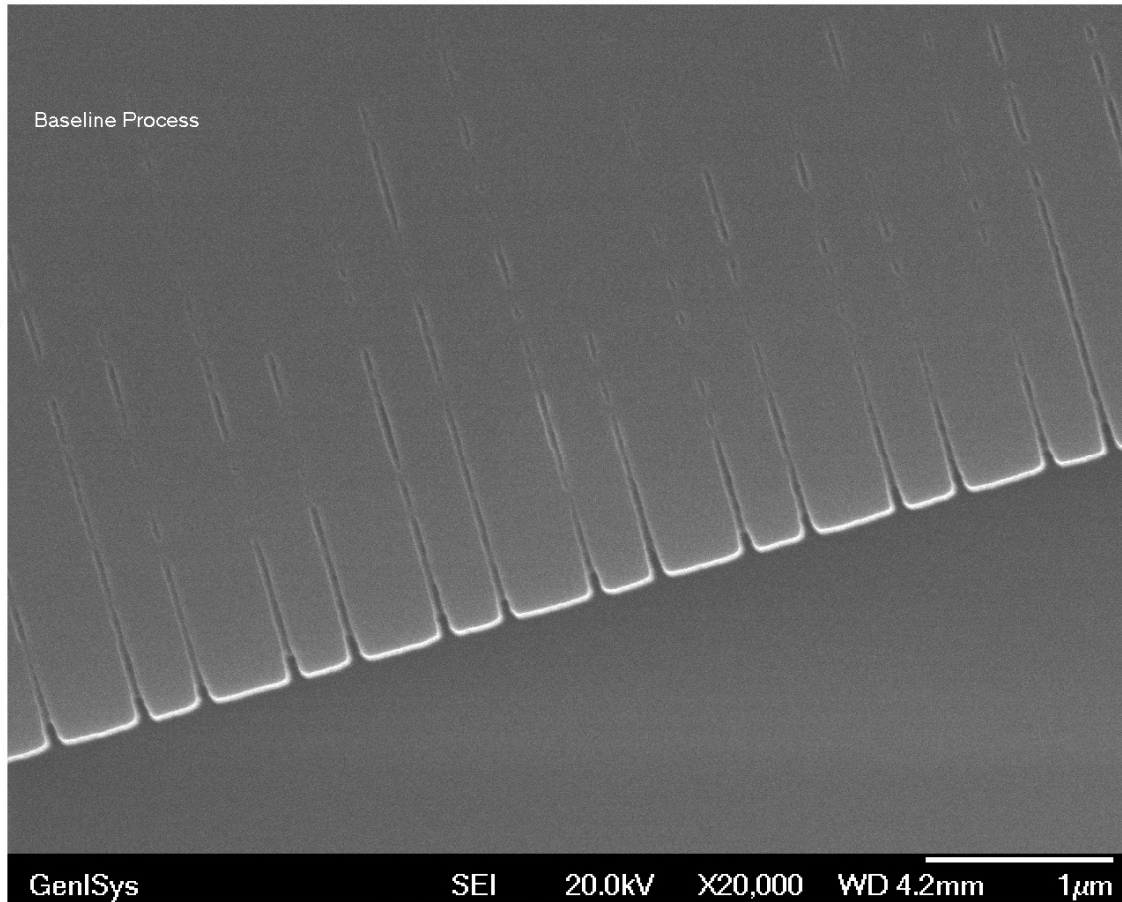
- Part 3 Summary: Dose PEC Parameter
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HSQ Silicon Photonics

- An HSQ process for Silicon Photonics has been in use for 9+ years.
- The process point was determined in a “traditional” way.
 - Use a baseline PSF for 100 kV electrons on Si.
 - Expose a dose matrix of the patterns, which were low-density waveguides (0-25%)
 - Choose base dose by observation, what dose gives proper size for a waveguide
- Hundreds of successful wafers have been built with this process.



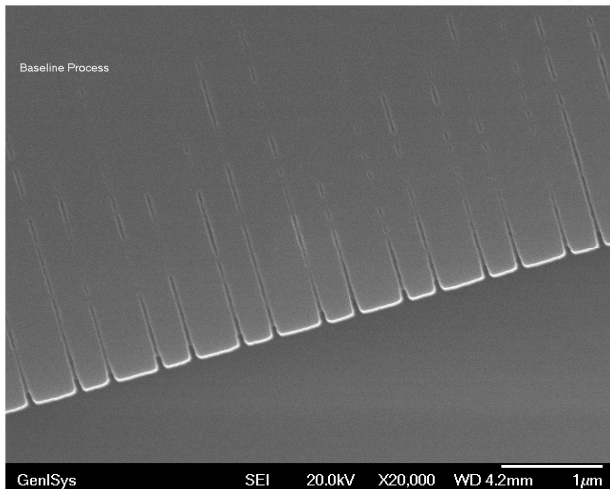
... until they needed higher-density patterns



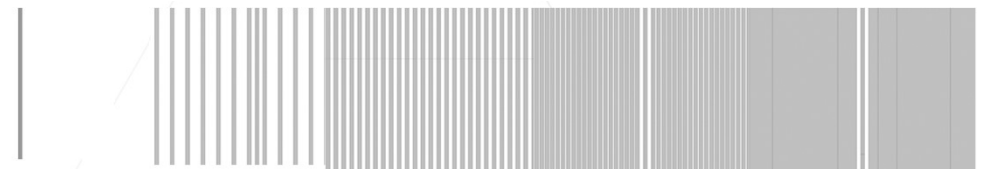
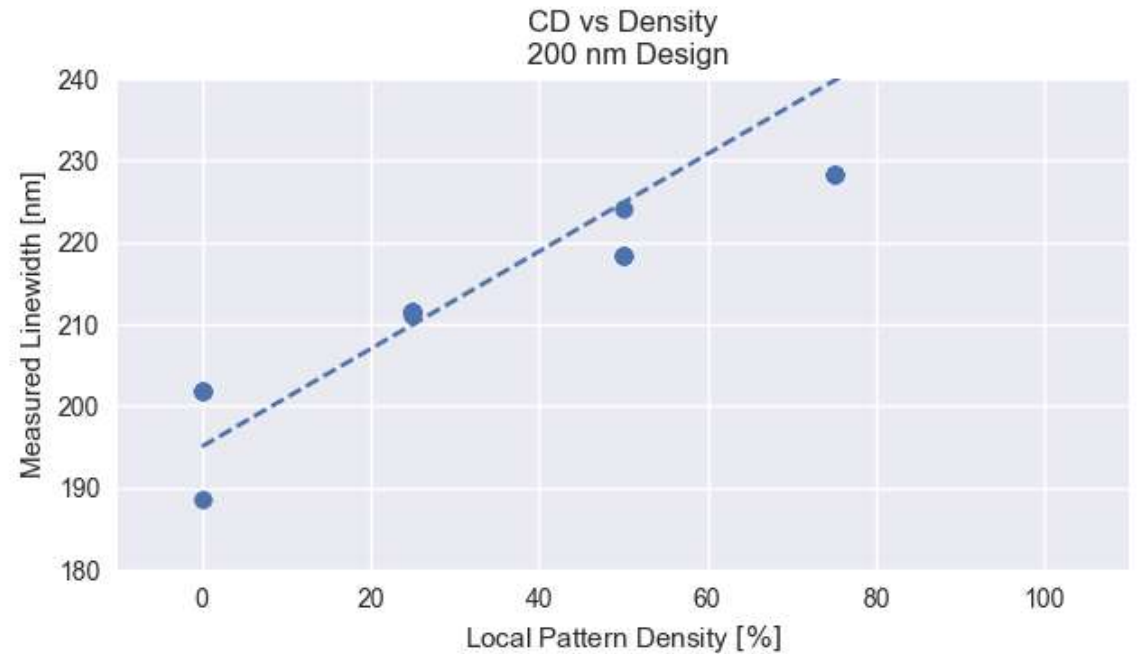
“Giant” subwavelength surface grating coupler.

Local density in the middle \sim 82%

- Strong dependence of linewidth on pattern density, even using baseline proximity effect correction.
- Dense patterns with small spaces are impossible.

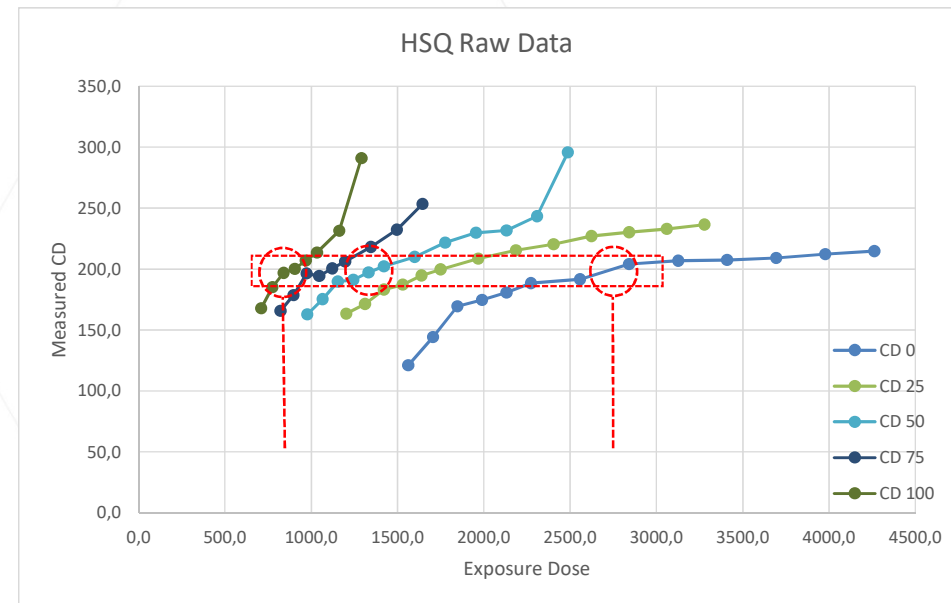
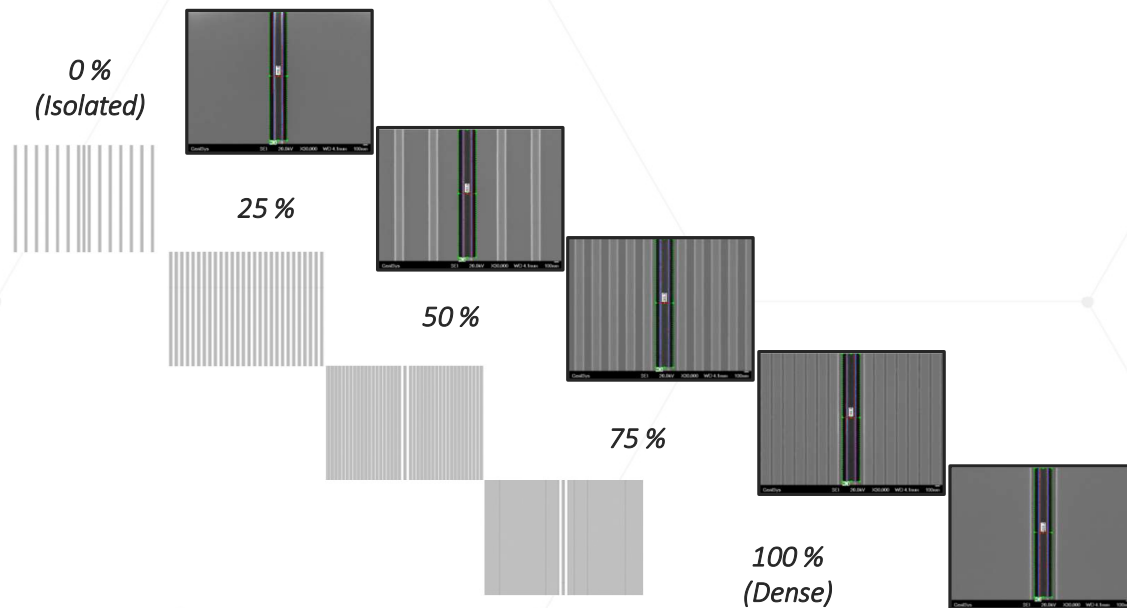


CD versus Local Pattern Density



Measuring Effect of Pattern Density

- Expose lines over a range of doses, with local pattern density varying from isolated to fully dense.



**Indicator: dose ratio D_{iso} / D_{dense} highly unusual: $2800 / 850 = 3.3$ (expected would be 2.2)
→ something in addition to scattering effects**

What is 'special' about HSQ?

- Many researchers have reported exposure effects such as neighboring-shape interactions beyond electron scattering, non-reciprocity (writing order) effects, or have observed the utility of adding an additional midrange Gaussian energy term to the dose-based PEC to improve results.
- Examples include: Liddle (2003), Olynick (2006), Brown (2013)
- Olynick 2006 speculates these are due to diffusion of hydrogen released during the exposure, which increases the HSQ sensitivity (lower the dose) of nearby shapes.
- However, no known published description of systematically quantifying and correcting for these effects.

Nanoscale topography control for the fabrication of advanced diffractive optics

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Scanning x-ray microscopy investigations into the electron-beam exposure mechanism of hydrogen silsesquioxane resists

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Lawrence Berkeley National Laboratory, Chemical Sciences Division, Berkeley, California 94720;
Department of Chemistry, University of California, Berkeley, California 94720; and Department of Physics,
University of California, Berkeley, California 94720

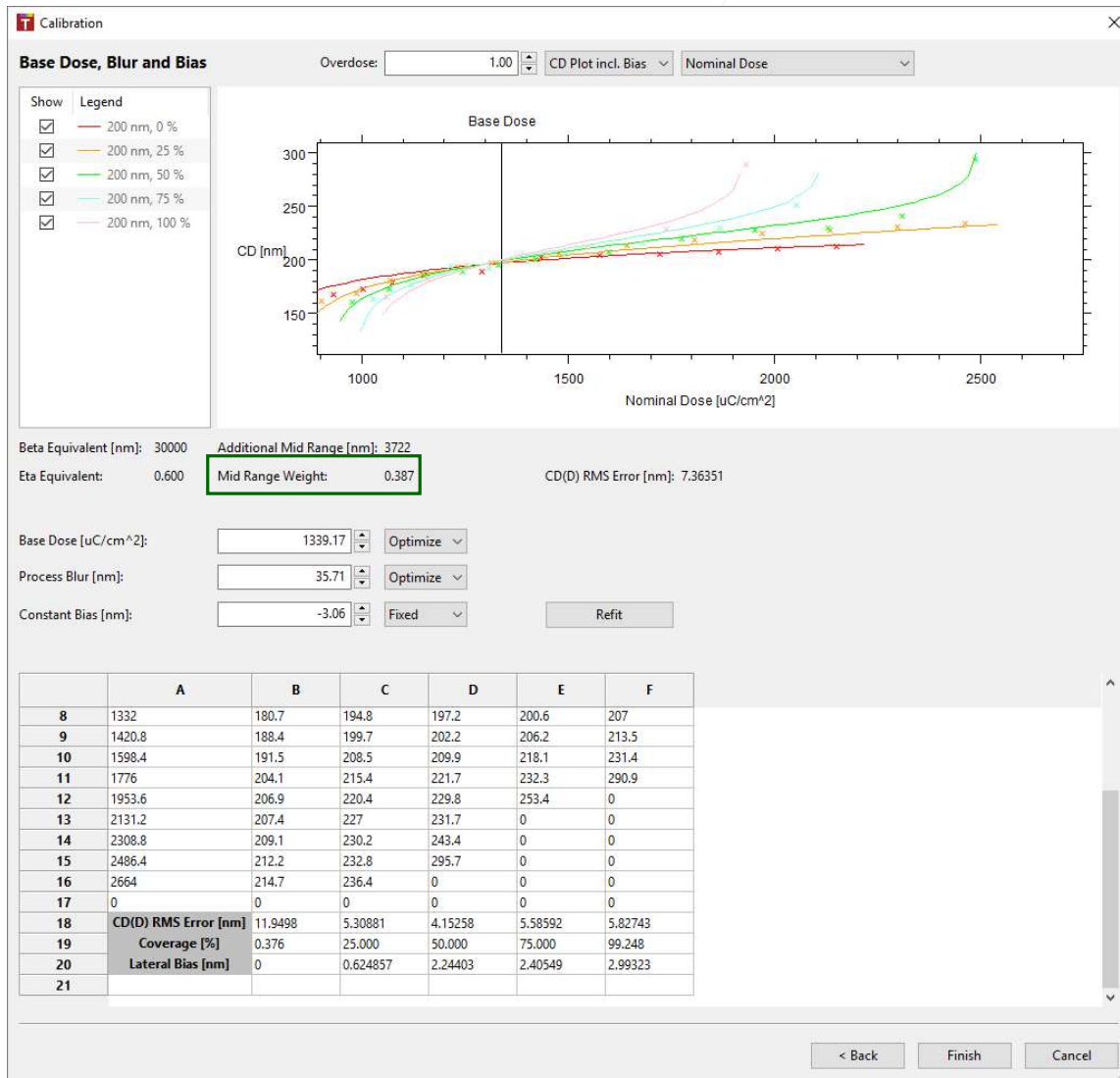
(Received 29 August 2006; accepted 18 October 2006; published 30 November 2006)

Pattern exposure order dependence in hydrogen silsesquioxane

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Hydrogen diffusion can be modeled by additional mid-range Gaussian

Fit Results



- The fitting procedure results in an “Extended Point Spread Function”, adding terms to the scattering PSF

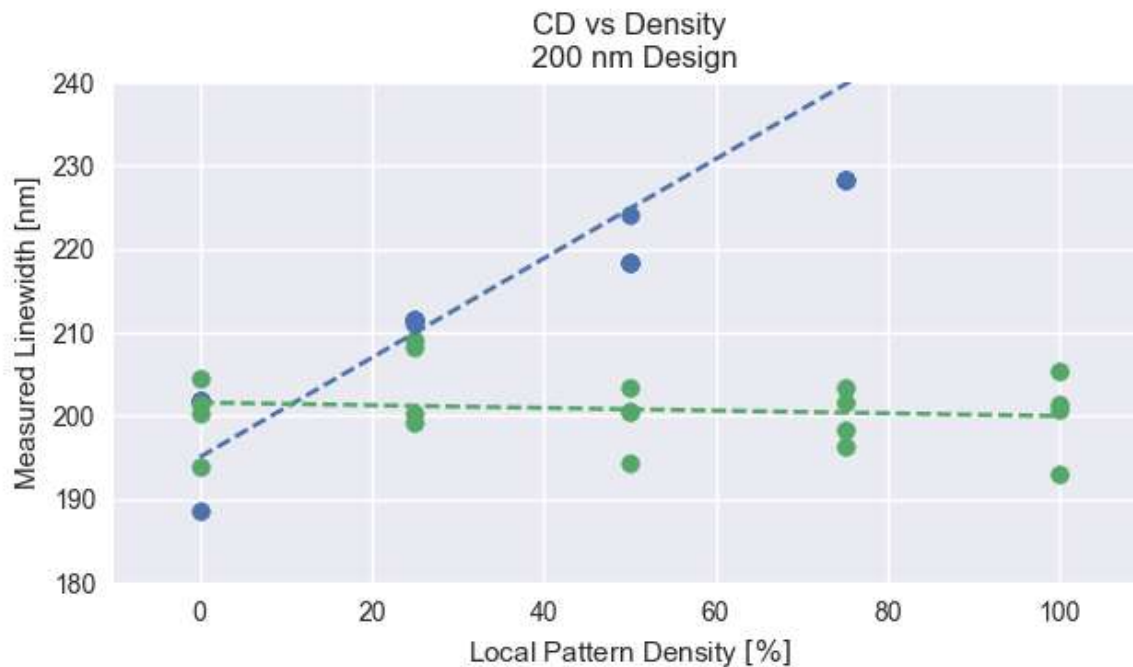
- Additional Midrange Gaussian term to compensate for hydrogen diffusion effects

- Overall Process Bias
- Density-dependent Bias terms to compensate for lateral development
- Optimal Base Exposure Dose
- Process Blur term

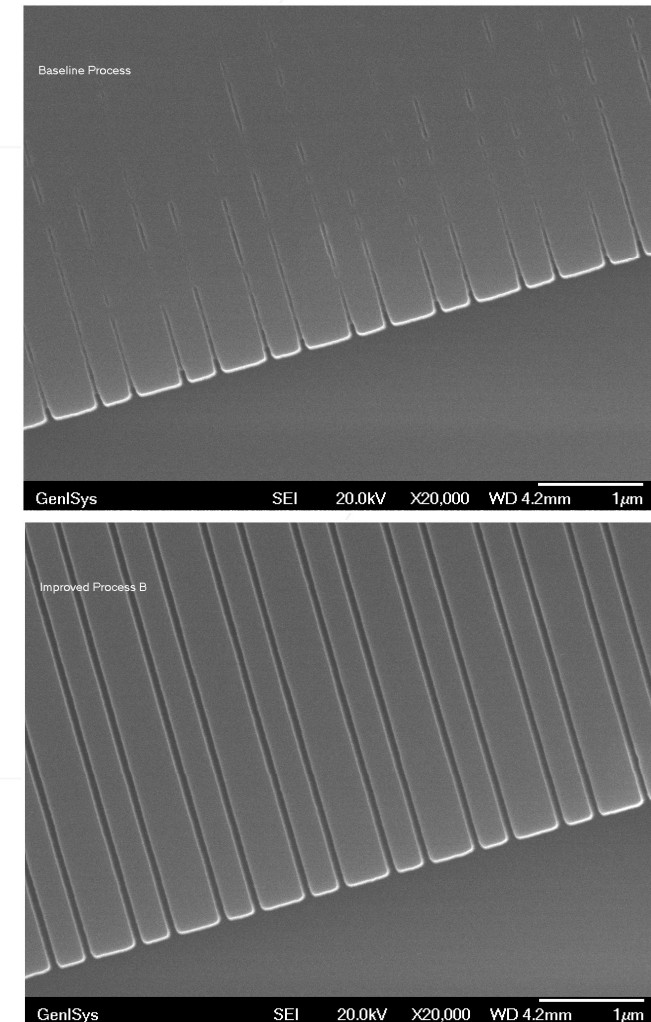
Extended PSF Calibration

	Before Calibration Chosen by 'Traditional Method'	Calibration Parameters Determined by TRACER fit to Measured CD Data
Base Dose	1750 $\mu\text{C}/\text{cm}^2$	1340 $\mu\text{C}/\text{cm}^2$
Process Blur	50 nm	36 nm
Process Bias	None	-3 nm
Density-dependent Bias	None	0% = 0 nm, 25% = 1 nm, 50% = 2 nm, 75% = 2 nm, 100% = 3 nm
PEC Parameters	Standard Si PSF	Additional Mid-range Gaussian $\gamma = 3722^* \text{ nm}$, $\nu = 0.38^*$

- With the enhanced correction:
 - Features are nominally the design size through the full range of density
 - Devices not previously successful with standard PEC correction are now fabricated successfully



Success



HSQ Summary

- HSQ exposures show (among other challenges):
 - Significant additional proximity effect, where exposed shapes can be affected by prior exposure of nearby shapes, above what is explainable by electron scattering
 - The effective dose for exposure is lowered when nearby shapes have been exposed
 - This effect can be measured and compensated for using dose proximity-effect correction, by treating the neighborhood exposures as an additional mid-range proximity effect term
- Not discussed here, but still true:
 - This also causes a significant “write-order” effect, so the sequence in which nearby shapes are written affects the resulting dimension
 - This write-order effect can be mostly mitigated by using multi-pass writing strategy

- Part 3 Summary: Dose PEC Parameter
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- “Real” processes have many effects beyond electron scattering
 - Process / metrology bias
 - Lateral development from finite resist contrast (density dependent)
 - Additional midrange process effects
- PEC Dose Range depends on
 - Resist contrast: consequence of the iso-focal shift (image iso-focal -> process iso-focal)
 - High-contrast requires $D_{iso}/D_{dense} = 1 + 2*BS/FS$, one PMMA required $D_{iso}/D_{dense} = 1 + 1.2*BS/FS$
 - Additional terms such as resist sensitivity changes (e.g. coming from catalytic reactions)
- TRACER can plot and fit the experimental data, providing the necessary process correction parameters
 - Maximizing the blur latitude to minimize process variation, e.g. across field
 - May include mix factor strategies, between Optimum Contrast and Uniform Clearing
 - Substrate and contrast dependent
 - Ability to adjust with parameters & see effects on process window, e.g. Undersize/Overdose
- BEAMER can be used to correct for not only the “Proximity Effect” but also these additional process effects

- Considers contrast curve
- Supports OC / UC
- Fit Additional mid-range term
- Fit „mix factor“

Calibration

Data

Name:

Description:

Preconditions for the TRACER Calibration include:
 1. An analytic PSF or a PSF from the archive
 2. A Dose vs. Density table obtained by exposing and evaluating a PEC corrected density varying pattern, obtainable from GenISys.
 3. Resist contrast value.

PEC Parameter used to process the calibration pattern

Use analytical PSF
 Beta [nm]: Eta:
 Gamma [nm]: Nu:

Use PSF from archive
 2D-PSF: Archive...

Optimal contrast [%]: / : Uniform clearing [%]

Calibrated Model

Resist: Resist contrast: Thickness [nm]: D0 [uC/cm^2]: From CC...

Use additional Mid Range Fit term
 Fit mix Factor
 Favor low-density fit quality

Separate low-density data fit [%]:

	A	B
1	Target CD [nm]	0
2	Density [%]	0.000
3	Dose [uC/cm^2]	Mea. CD [nm]
4	0	0

Add Dose
 Add Dataset
 Remove
 Import...
 Export...

< Back Next > Cancel