ICTP winter college 7-18 February 2005 Trieste **Introduction to nanofabrication** LILIT beamline INFM (National Institute for Matter Physics) @ Elettra Synchrothron Light Source -S.S 14 Km 163.5, Area Science Park, 34012 Basovizza- Trieste (Italy) **L.I.LIT** *Laboratory for Interdisciplanry LIThography Enzo Di Fabrizio*

Lesson plan

- Lesson 1: introduction to nanofabrication
- Lesson 2: diffractive optics and X-ray microscopy
- Lesson 3: 3D fabrication and optical manipulation and spectrosopy

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TASC National Laboratory of INFM

Tools to observe the nano world

• (SPM, scanning probe microscope):

STM (scanning tunneling microscope) Resolution ∼0.1 nm AFM (atomic force microscope) Resolution ∼1 0nm

- SEM (scanning electron microscope) Resolution ∼1 nm
- TEM (transmission electron microscope) Resolution ∼0.1 nm

STM microscope

Muovendo la punta sopra la superficie, si misura la corrente di elettroni che che passano dallla superficie alla punta (o viceversa) per effetto tunnel. Spostando la punta con dei piezoelettrici (precisione 0.01 nm) si ottiene una mappa della densita' elettronica.

> Densita' elettronica di una superficie di Nickel (110). Realizzazione ed elaborazione IBM

Immagine di una punta per STM

AFM microscope

Muovendo la punta sopra il campione, si misura il piegamento della piccola leva (canti-lever) dovuto alle forze di interazione atomiche e molecolari.

Si ottiene una immagine topografica della superficie che consente di misurare anche materiali isolanti (organici, biologici)

Immagine di una punta montata su una piccola leva per AFM

- **Fundamentals of Lithography**
- **Beam Lithography**
- **Alternate Nanolithography Techniques**

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The Si revolution…

First Transistor

Bell Labs (1947) Si integrated circuits Texas Instruments (~1960)

Modern ICs

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The need of micropatterning

The batch fabrication of microstructures requires a low-cost, high throughput surface patterning technology

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Elements of photolithography

- **Lithography consists of patterning a substrate by employing the interaction of beams of photons or particles with materials.**
- **Photolithography** is widely used in the integrated **circuits (ICs) manufacturing.**
- **The process of IC manufacturing consists of a series of 10-30 steps or more, called mask layers where layers of materials coated with resists are patterned then transferred onto the material layer.**

Output Spectrum of Hg Arc Lamp

Size Scales Accessible to Nanofabrication Approach

Elements of photolithography (ctnd.)

- **A photolithography system consists of a light source, a mask, and a optical projection system.**
- **Photoresists are radiation sensitive materials that usually consist of a photo-sensitive compound, a polymeric backbone, and a solvent.**
- **Resists can be classified upon their solubility after exposure into: positive resists (solubility of exposed area increases) and negative resists (solubility of exposed area decreases).**

Micro- and nanolithography

- **Present techniques in IC manufacturing involversion** dimensions in order of 100-150 nm.
- Diffraction and other optical effects limit the resolution of "standard" UV photolithography to the ~100 nm range.
- Photolithography continues to support IC manufacturing in the sub-100 nm region through continuous advances in optics (UV to DUV to EUV) and resist engineering.
- Exploratory research in the sub-100 nm region may also be accomplished through alternate patterning techniques such as *x-ray-*, *ion-* and *electron beam*lithography.

BEAM ASSISTED LITHOGRAPHY

- **X-ray**
- **Electron Beam**
- **Ion Beam**

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X Rays

Features:

- **Discovered by Wilhelm Conrad Röntgen in 1895**
- **Experiments in a vacuum tube made a nearby fluorescent screen glow.**

1901 W. C. Roentgen in Physics for the discovery of x-rays.

- 1914 M. von Laue in Physics for x-ray diffraction from crystals.
- 1915 W. H. Bragg and W. L. Bragg in Physics for crystal structure derived from x-ray diffraction.
- 1917 C. G. Barkla in Physics for characteristic radiation of elements.
- 1924 K. M. G. Siegbahn in Physics for x-ray spectroscopy
- 1927 A. H. Compton in Physics for scattering of x-rays by electrons.

1936 P. Debye in Chemistry for diffraction of x-rays and electrons in gases.

- 1962 M. Perutz and J. Kendrew in Chemistry for the structure of hemoglobin.
- 1962 J. Watson, M. Wilkins, and F. Crick in Medicine for the structure of DNA.
- 1979 A. McLeod Cormack and G. Newbold Hounsfield in Medicine for computed axial tomography.
- 1981 K. M. Siegbahn in Physics for high resolution electron spectroscopy.
- 1985 H. Hauptman and J. Karle in Chemistry for direct methods to determine x-ray structures.

1988 J. Deisenhofer, R. Huber, and H. Michel in Chemistry for the structures of proteins that are crucial to photosynthesis.

THEORETICAL UNDERSTANDING \rightarrow

1873 **Maxwell's equations**

 \rightarrow made evident that changing charge densities would result in electric fields that would radiate outward

Heinrich Hertz demonstrated such waves: 1887

..... this is of no use whatsoever !

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Why do they radiate?

Charge at rest: Coulomb field

Uniformly moving charge

Accelerated charge

Fields of a moving charge

$$
\vec{\mathbf{E}}(t) = \frac{q}{4\pi\epsilon_0} \left[\frac{\vec{\mathbf{n}} - \vec{\beta}}{\left(1 - \vec{\mathbf{n}} \cdot \vec{\beta}\right)^3 \gamma^2} \cdot \frac{\left[1}{r^2}\right]_{ret} + \cdots \right]
$$

$$
\frac{q}{4\pi\epsilon_0 c} \left[\frac{\vec{\mathbf{n}} \times \left[(\vec{\mathbf{n}} - \vec{\beta}) \times \vec{\beta} \right] \cdot \left[\frac{1}{r} \right] \right]_{ret}
$$

$$
\vec{\mathbf{B}}(t) = \frac{1}{c} [\vec{\mathbf{n}} \times \vec{\mathbf{E}}]
$$

Transverse acceleration

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Radiation field quickly
separates itself from the
Coulomb field

Longitudinal acceleration

Radiation is emitted into a narrow cone

Into a narrow cone

Crab Nebula 6000 light years away

First light observed

1054 AD

GE Synchrotron New York State

First light observed 1947

20 000 users world-wide

Scheme of a typical synchrotron X-ray lithography system

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The Stepper

11

The absorption from two Berillium windows of various thicknesses is included in the calculation.

Layered approach for the E.M. field

Figure 8: Illustration of the layered approach. Materials of arbitrary shape can be treated as a stack of layers with varying transmission. The figure is not drawn to scale, in order to show the structure clearly. A spherical particle in the mask can be approximated by circular discs

ICTP 6: go to next layer **2005** Triester college 7-18 February 2005 Triester 2005 Triester 2005 Triester 2005 **Step 1**:Compute the transmission of the l-th layer **Step 2**:Compute the transmitted E.M. field **Step 3**: Obtain the Fourier Tranf. of the E.M.field **Step 4**: Propagate the field through the thickness d in the Fourier domain **Step 5**:obtain the diffracted E.M. field by inverse FFT

1:1 XRL for 2D GaAs/AlGaAs PC

X-ray Replication X-ray Mask & Liftoff (Ti/Au) 33 nm 33 nm

Device patterns with feature sizes less than 40 nm achieved by xray lithography and by liftoff.

> ICTP winter college 7-18 February 2005 Trieste **More ? Check out Prof. H. I. Smith, MIT**

Structures produced with X-ray litho. (ctnd.)

200 nm lines of 2 um thick resist over thick steps (NTT-AT)

ICTP winter college 7-18 February 2005 Trieste **400 nm by 3 um resist over poly line (ISiT)**

Source: SAL, Inc.

Electron-Beam nanolithography syetsm Installed in Pirelli Labs-Milan Electron-Beam nanolithography syetsm Installed in Pirelli Labs-Milan

Developed further to the needs of micro-electronics industry, electron-beams are systems able to impress a thin layer of photosensitive material (photoresist) with a resolution orders of magnitude higher than that achieved by optical methods

Typical parameters of operation:

- •spot size of the beam: 5nm
- •resolution: 15nm (conservative)

•size of the wafer: 8", 12" standard, larger sizes developed for semiconductor industry

Typical e-beam system used in the production (Leica VB6 HR type)

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Electron beam lithography system

Effects produced by electron bombardment of a material.

Two major factors control which effects can be detected from the interaction

volume. First, some effects are not produced from certain parts of the interaction volume (Figure 2.1b).

Beam electrons lose energy as they traverse the sample due to interactions with it and if too much energy is required to produce an effect, it will not be possible to produce it from deeper portions of the volume. Second, the degree to which an effect, once produced, can be observed is controlled by how strongly it is diminished by absorption and scattering in the sample.

For example, although secondary and Auger electrons are produced throughout the interaction volume, they have very low energies and can only escape from a thin layer near the sample's surface. Similarly, soft X-rays, which are absorbed more easily than hard X-rays, will escape more readily from the upper portions of the interaction volume. Absorption is an important phenomenon and is discussed in more detail below.

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Figure 2.1b. Generalized illustration of interaction volumes for various electron-specimen interactions. Auger electrons (not shown) emerge from a very thin region of the sample surface (maximum depth about 50 Å) than do secondary electrons (50-500 Å).

Volume of Excitation

Two factors limit the size and shape of the interaction volume: (1) energy loss through inelastic interactions and (2) electron loss or backscattering through elastic interactions. The resulting excitation volume is a hemispherical to jugshaped region with the neck of jug at the specimen surface. The analyst must remember that the interaction volume penetrates a significant depth into the sample and avoid edges where it may penetrate overlapping materials. The depth of electron penetration of an electron beam and the volume of sample with which it interacts are a function of its angle of incidence, the magnitude of its current, the accelerating voltage, and the average atomic number (Z) of the sample. Of these, accelerating voltage and density play the largest roles in determining the depth of electron interaction (Figure 2.2a).

Figure 2.2a. Schematic depiction of the variation of interaction volume shape with average sample atomic number (Z) and electron beam accelerating voltage (E_0) . The actual shape of the interaction volume is not as longnecked since the electron beam in microprobe analysis has a diameter of about 1 μ m (see Figure 2.1b).

$$
x \, (\mu m) = \frac{0.1 \, E_o^{1.5}}{\rho}
$$

where E_0 = accelerating voltage (keV), and $\rho =$ density (g/cm³)

Electron penetration generally ranges from 1-5 μ m with the beam incident perpendicular to the sample. The depth of electron penetration is approximately (Potts, 1987, p. 336):

For example, bombarding a material with a density of 2.5 g/cm3, about the minimum density for silicate minerals, with $E_0 = 15$ keV, gives $x = 2.3$ µm. The width of the excited volume can be approximated by (Potts, 1987, p. 337):

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Both of these are empirical expressions. A theoretical expression for the "range" of an electron, the straight line distance between where an electron enters and its final resting place, for a given E_{α} is (Kanaya & Okayama, 1972):

The volume of interaction can be modeled by Monte Carlo simulation. In such models, the likelihood of incident electrons interacting with the sample and scattering and the angle of deflection are determined probabilistically. X-ray generation depths depend strongly on density and accelerating voltage (Figure 2.2b.). The results derived from Monte Carlo modeling yield a volume of interaction that is very similar to that determined by etching experiments. The excited volume is roughly spherical and truncated by the specimen surface. The depth of the center of the sphere decreases with increasing atomic number of the target.

$$
y \text{ (µm)} = \frac{0.077 \text{ E}_o^{1.5}}{\rho}
$$

where E_0 = accelerating voltage (keV), and $\rho =$ density (g/cm³)

$$
r \text{ } (\mu \text{m}) = \frac{2.76 \times 10^{-2} \text{ A} \text{ E}_0^{1.67}}{\rho \text{ } Z^{0.89}}
$$

where ρ = density of the material (g/cm^3), $Z =$ atomic number, $A = atomic mass$, and E_0 = accelerating voltage.

Figure 2.2b. Comparison of electron paths (top) and sites of X-ray excitation (bottom) in targets of aluminum, copper, and gold at 20 keV, simulated in a Monte Carlo procedure (after Heinrich, 1981).

PEC(Proximity Effect Correction) a. Dose Distribution due to PE

G (x) EID (Exposure Intensity Distribution)function (Dose distribution of EB resist at one point (x=0) irradiation)

$$
g(\mathbf{x}) = \mathbf{C} \left\{ g_f(\mathbf{x}) + \eta_E g_b(\mathbf{x}) \right\},\tag{1}
$$

$$
g_f(\mathbf{x}) = \frac{1}{\pi} \sigma_f^2 \exp\left\{-\left(\mathbf{x} - \mathbf{x}'\right)^2 / \sigma_f^2\right\} \qquad (2)
$$

$$
g_b(\mathbf{x}) = (1/\pi \sigma_b^2) exp \left(-\left(\mathbf{x} - \mathbf{x}'\right)^2 / \sigma_b^2\right)
$$
 (3)

(Parameter)

(Factor for parameter)

- **1. Material of substrate**
- **2. Acceleration voltage**
- **3. Resist material**
- **4. Resist thickness**

Simulation for Incident Electron trajectories

Vacc:50kV, **Substrate: Si Wafer**, **Resist Thickness**: **300nm**

Resist + Substrate Resist Part Enlargement

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Simulation for Incident Electron trajectory

Vacc:30kV, **Substrate:Si Wafer**, **Resist Thickness**: **300nm**

Resist + Substrate Resist Part Enlargement

Dose Distribution (DD) Simulation

* **Resist dose amount at X in case of irradiating pattern A**

 $E(x)=\int D(x')g(x-x')dx'$ **X**Э**A**

When,

Integration: area A irradiated by EB

D(x´**): Irradiation amount at x** ´

*** Pattern dependence of resist dose amount**

pattern density.

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Dose Distribution (DD) Simulation (2)

■ E B uniform irradiation

■ : E B uniform irradiation

DD of a,b,c change due to EB irradiation at d.

DD of a,b,c change again due to EB irradiation at e.

Dose Distribution Simulation vs Writing Result

(Simulation parameter) Backward scattering coeffi. : 3μ**m**

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Main PEC Methods

Proximity Effect for 100nm L&S Lattice

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PEC(Proximity Effect Correction) for 100nm L&S Lattice

(Writing Conditions) (PEC Conditions) Vacc: 30kV PEC: Irradiation amount correction Field Size: 500µm Correction Amount: 5 stage corr. as base of nominal dose on basis of simulation result Writing Area: 10 µm Resist / Thickness: ZEP520/300nm Nominal Dose: 50μ**C/cm2 (6.25**μ**sec/dot) Pixel: 20,000 x 20,000 dot I B: 50pA**

Resist Thickness Dependence of Proximity Effect

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Without PE Correction

Pattern design: 500mm field, 20000dot =Line 125nm/Space 75nm Size: 50mm x10mm

ZEP520 300nm thickness on Si sub.

 e^- Beam = 30kV / 50pA

With PEC Correction Collection point 2, connection 2/3, quantization level 6

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Proximity Effect Correction

(Data supplied by Thales Research & Technology)

Proximity Effect Correction

(Data supplied by Thales Research & Technology)

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L- SE1 EHT- 18.0 KV HD- 3
20.0um
C.N.R. I.E.S.S. MAG- X 922.

With proximity correction Without proximity correction

From the remaining resist thickness vs. absorbed dosage curve it is possible to interpolate the electron dosage values necessary to achieve determined resist thicknesses

Resist pads are exposed at increasing electron dosages (the pad lateral size is larger than the electron backscattering range \implies the exposure dosage is equal to the absorbed dosage)

CNR-IESS - µ**Fab Group**

Continuous profiling

Normalized Resist Thickness vs. absorbed dosage curve obtained with **MiBK-IPA 3:1**

Scanning Electron Microscope analysis SEM image of a 16-level, 32 steps double ramp-shaped pattern. Development: MiBK-IPA 3:1 for 20 sec. 16-levels Testing the dose calibration on a Expected profile $\begin{bmatrix} 1 & 45 \text{ nm} \\ 1.45 \text{ nm} \end{bmatrix}$ practically realized multi-level profile **CNR-IESS -** µ**Fab Group** MAG- X 21.5 K PHOTO- 10 $L - SE1$ T-18.0 KV WD-10 mm Profilometer measurements of the resist thickness CNR -0.5 performed on a 16-level -0.3 ramp-shaped pattern. -1.2 Development: MiBK-IPA -1.6 3:1 for 20 sec. The step width is 20 μm and their mean height is ∼100 nm. The maximum resist height is 1.5 μ m. **CNR-IESS -** µ**Fab Group**

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Applications of electron beam lithography

Mainly employed for the fabrication of photomasks

Also used to write patterns directly on wafer

L/S(Line & Space)Resist Pattern

HV : 30keV

Dose : 50μC/cm2

 $L/S = 50$ nm/70nm $L/S = 70$ nm/70nm

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L/S(Line & Space)Resist Pattern

HV : 50keV

Dose : 140μC/cm2

 $L = 50$ nm $P = 100$ nm

 $L = 70$ nm $P = 140$ nm

10 nm Space Width Resist Pattern 10 nm Space Width Resist Pattern

HV: 50kV Resist : ZEP520

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Fabrication of Multi-Pitch Gratings

Spot Scan Writing-Si Nanopillar for Photonic Crystal (by Spot Scan Writing)

 $\overline{\mathbf{o}}$

 $\overline{\circ}$

Nano-pillar **Dot array**

o

 $\ddot{\circ}$

 $\ddot{\circ}$

The data supplied by Dr.T,Kanayama and Dr.T.Tada of JRCAT

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Hexagonal Grating (by Spot Scan Writing)

500 dots/100μ**m- length**

30kV 5×**10-11A 40**μ**s/dot**

3-D Concentric Circular Pattern

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Continuous profile holographic surfaces produced by e-beam methods Continuous profile holographic surfaces produced by e-beam methods

Continuous profile make it possible to design a wavefront transforming surface having the exact theoretical shape required to maximize efficiency.

•*Continuous profile*

•*Overall pattern*

•*Uniform redistribution of laser light obtained by the innovative optical element*

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Ion beam lithography techniques

Focused Ion Beam Lithography (FIBL)

Masked Ion Beam Lithography (MIBL)

Ion Projection Lithography (IPL)

Focused Ion Beam Lithography **FIBL components**: ON SOURCE – Ion source – Ion optics column – Sample displacement table **Specifications:**

- Accelerating voltage 3-200 kV.
- Current density up to 10 $A/cm²$.
- Beam diameter 0.5-1.0 µm.
- Ions: Ga+ , Au+ ,Si+ ,Be+ etc.

FIB fabricated nanostructures

Ion projection lithography

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Alternate Nanolithography **Techniques**

-
-
-
-

Micro-contact printing

Printing of PDMS

High-resolution µCP of 60 nm dots

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NIL master

- SiO₂ pillars with 10 nm diameter, 40 nm spacing, and 60 nm height fabricated by e-beam lithography.
- Master can be used tens of times without degradation

NIL pattern in PMMA

- Mask is pressed into 80 nm thick layer of PMMA on Si substrate at 175° C $(\mathsf{T}_{q} = 105 \degree C)$, P= 4.4 MPa.
- PMMA conforms to master patterng, resulting in ~10 nm range holes

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Carbon nanotubes for nanolithography

- A carbon nanotube can be used as a tip in an atomic force microscope (AFM). Such a tip in an AFM can be used to create nanoscale patterns i.e. nanolithograpghy or to etch material away from a surface in the fabrication of semiconductor chips
- The videoclips show real-time dynamics of interaction between carbon nanotube tips and silicon and surfaces

