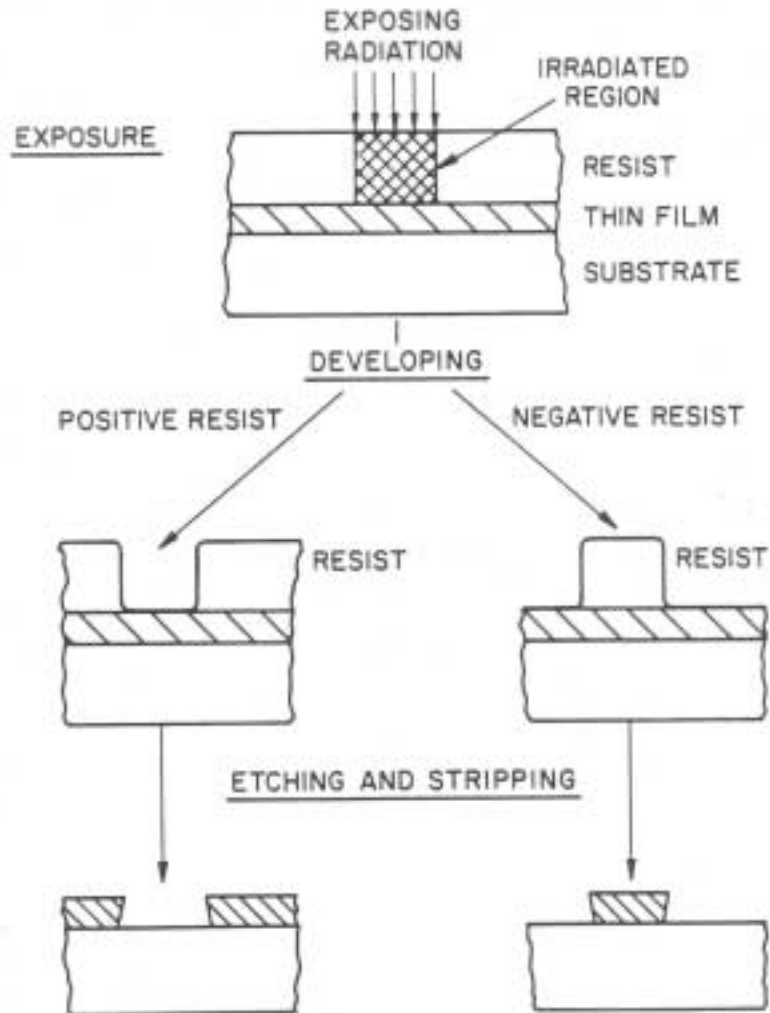


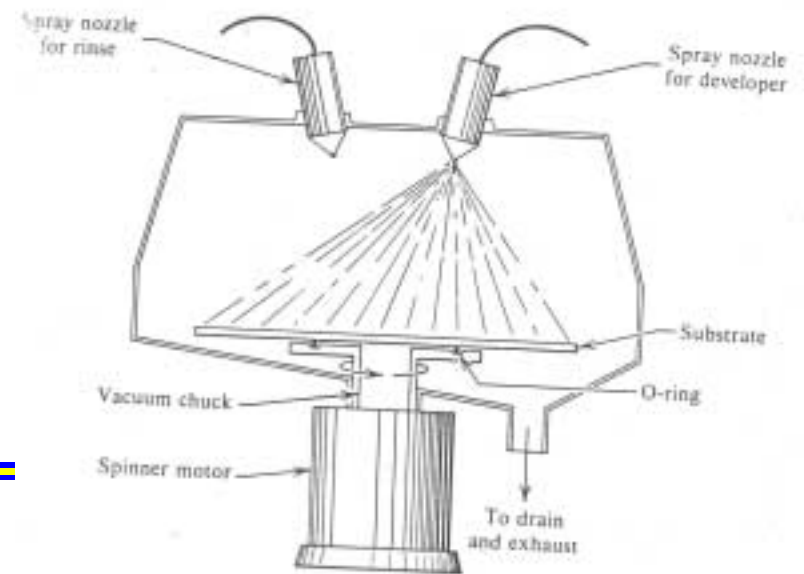
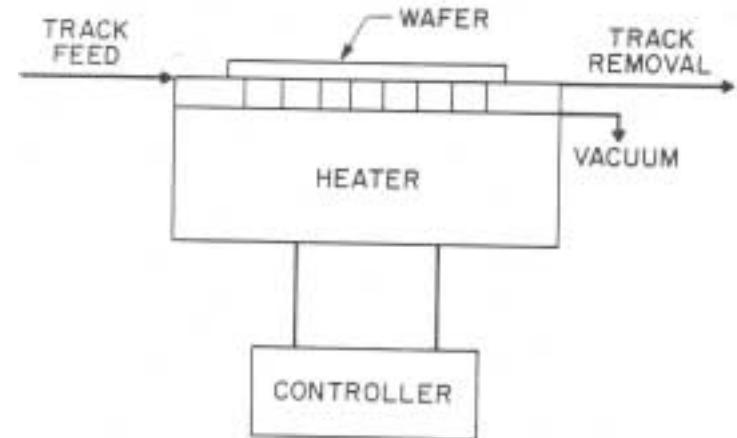
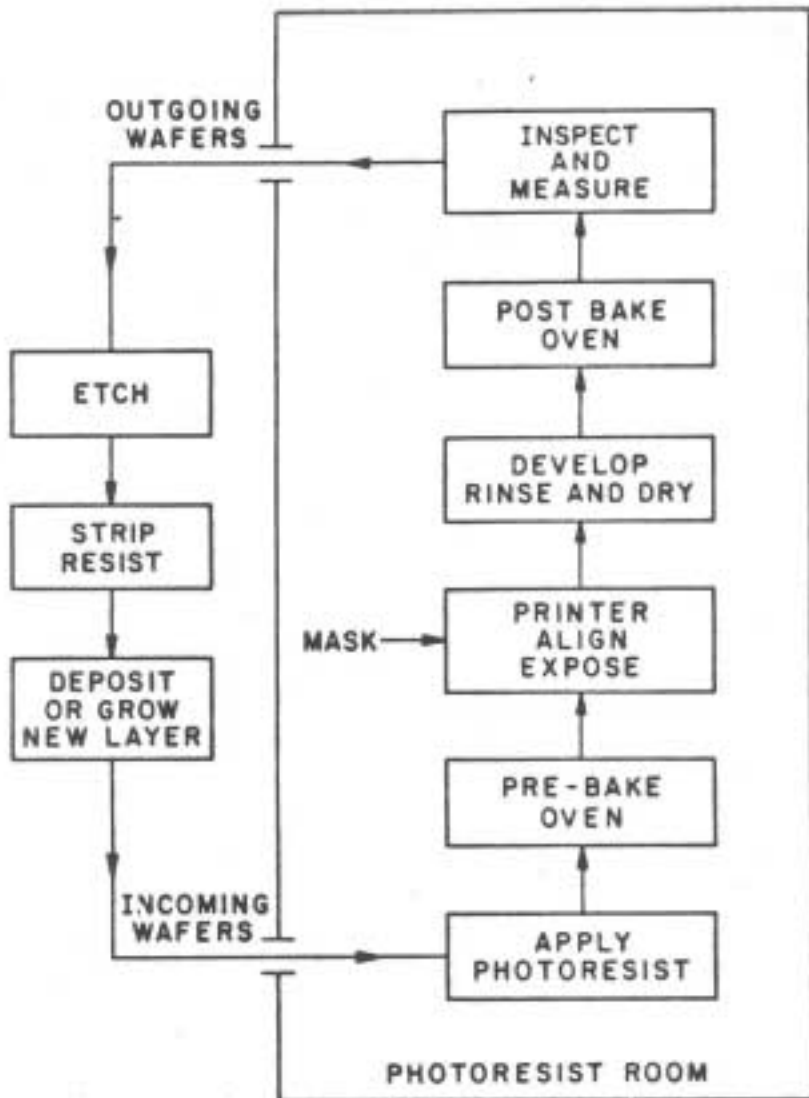
Lithography

- **Lithography** is the process of transferring two dimensional patterns (e.g., a circuit pattern) on a mask to an underlying film or substrate.
 - Most often the mask is a light sensitive polymer which allows the patterns to be transferred onto the polymer film. Such a light sensitive polymer is called a **photoresist**.
 - Photolithography uses ultraviolet radiation from a Hg light source (now we use excimer lasers) to transfer the pattern from the mask to the photoresist and is the dominant technology in IC fabrication today.
 - Future generations of lithography make use of x-rays, electrons or focused ion beams instead of photons (light).
-

Patterning thin films by Lithography



Lithography is used many times in IC manufacturing and complexity of the IC is measured in “mask levels”



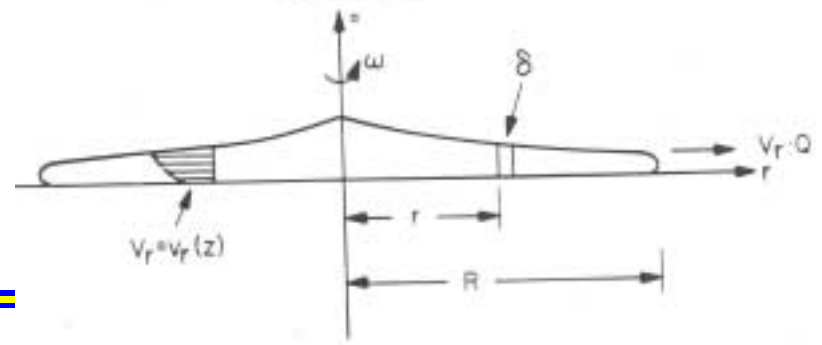
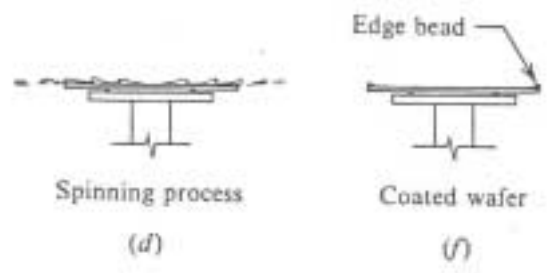
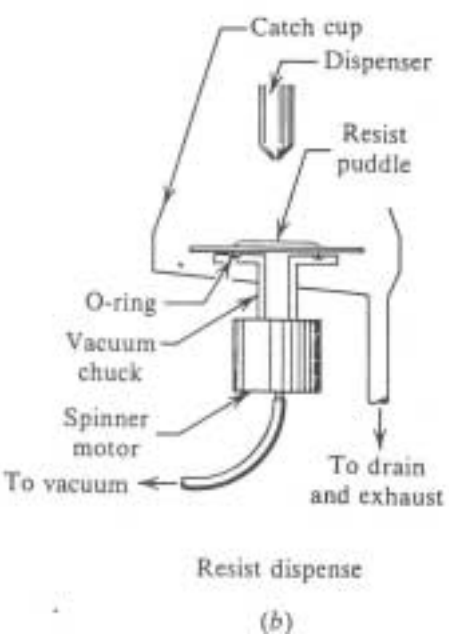
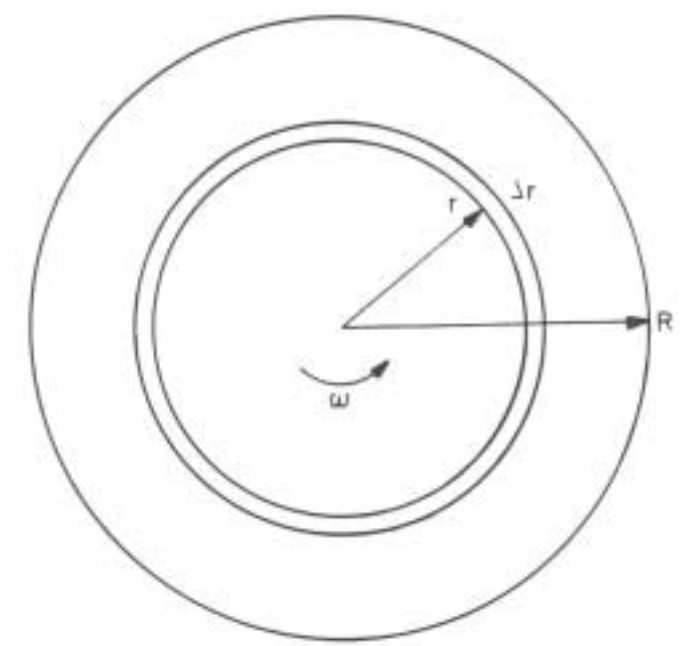
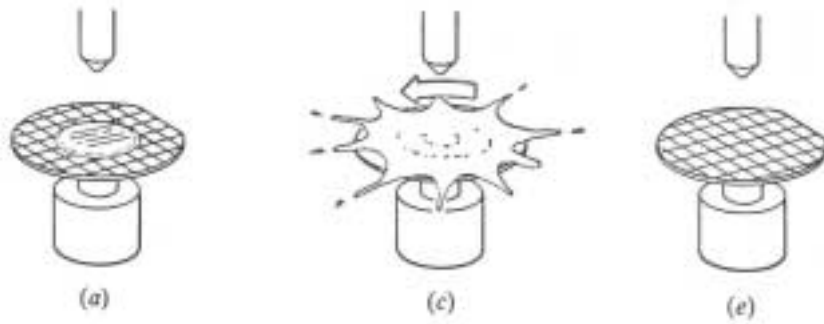
Application of photoresist

- Photoresist are high molecular weight polymers dissolved in organic solvents
- The goal in applying the photoresist is to obtain uniform, adherent, defect free polymeric film over the entire substrate. Uniformity must be ± 5 nm.
- The polymer solution is spin coated. Spin coating is accomplished by flooding the substrate surface with the resist solution and rapidly rotating it at constant speed. $\omega \sim 1000-10,000$ rpm.
- First the substrate is cleaned. Contaminated substrate surface is a source of problems. Contamination causes poor adhesion and particles cause “comet tail” defects.
- Polymer must be compatible with the substrate surface for good adhesion. Sometimes adhesion promoters such as Hexamethyldisiloxane (for Si substrates) are used.

Spin coating

- Consists of 4 steps
 - 1) Flooding the substrate with the resist solution.
 - 2) Accelerating to the desired rotation speed.
 - 3) Spinning at constant speed to near dryness.
 - 4) Remove the edge bead.
- The solvent evaporates as the substrate is spinning and the solution is spreading on the substrate.
- Thickness t is a function of ω, c, η ; $t = f(\omega, c, \eta, \text{solvent, environmental factors})$
 - ω = spin speed
 - c = polymer concentration in solvent
 - η = polymer viscosity

Photoresist Spinning



Spin coating

- For a given system of polymer and solvent $t = f(\omega)$ only.

$$t = \frac{k}{\omega^\alpha} \quad \alpha \approx 1/2$$

- If polymer concentration increases k is increased $k=k'c^\beta$.

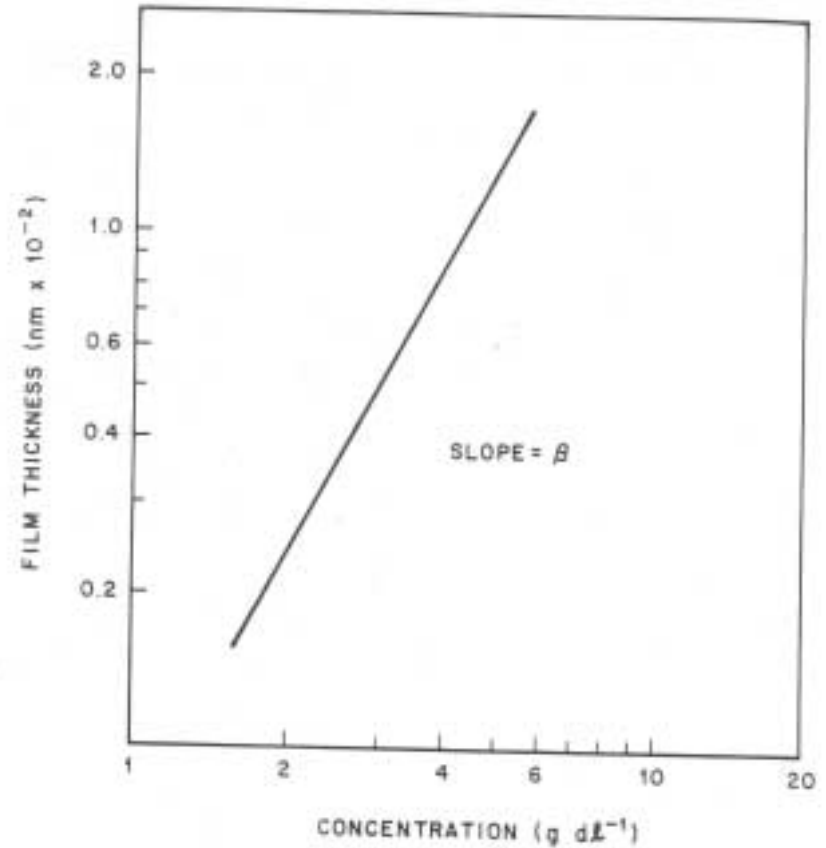
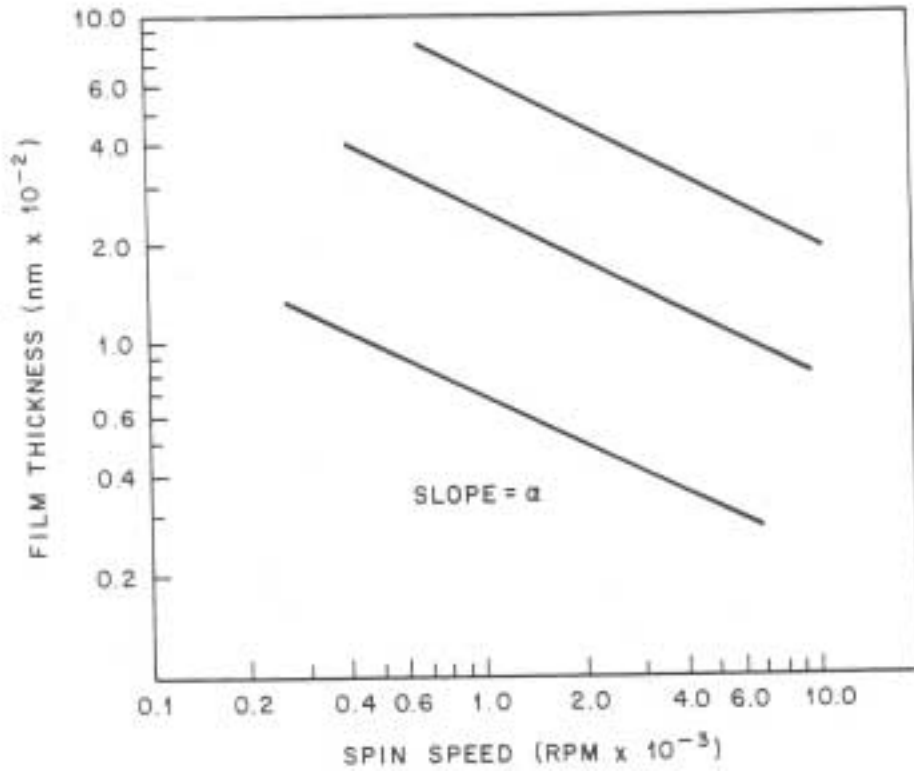
- Similarly s a function of viscosity k increases $k=k''\eta^\gamma$.

- Combining these gives

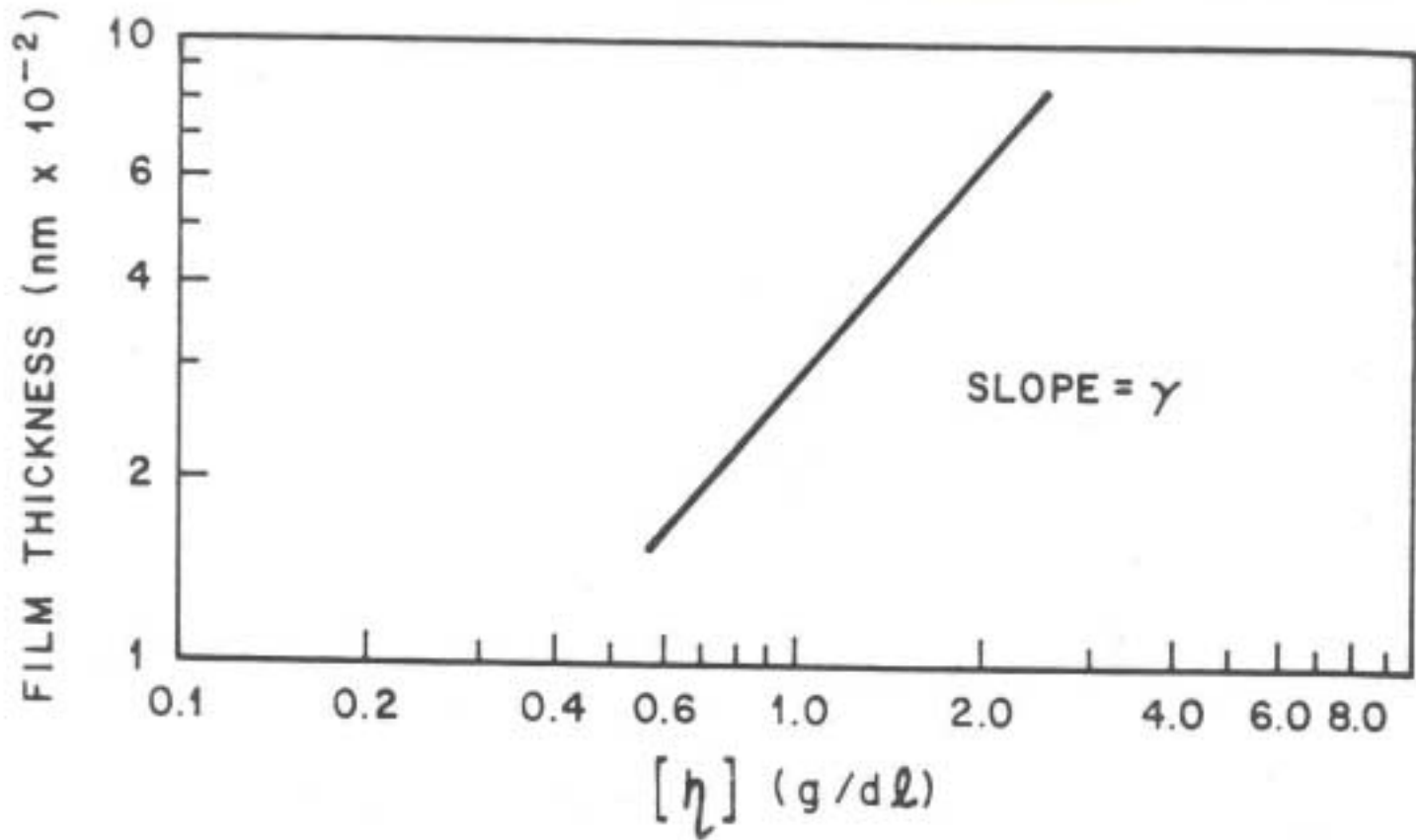
$$t = K \frac{c^\beta \eta^\gamma}{\omega^\alpha}$$

- For a given photoresist system this data would be supplied by the photoresist supplier.

Thickness t is a function of $\omega =$ spin speed
 $c =$ polymer concentration in solvent



Thickness as a function of the viscosity



Spin coating

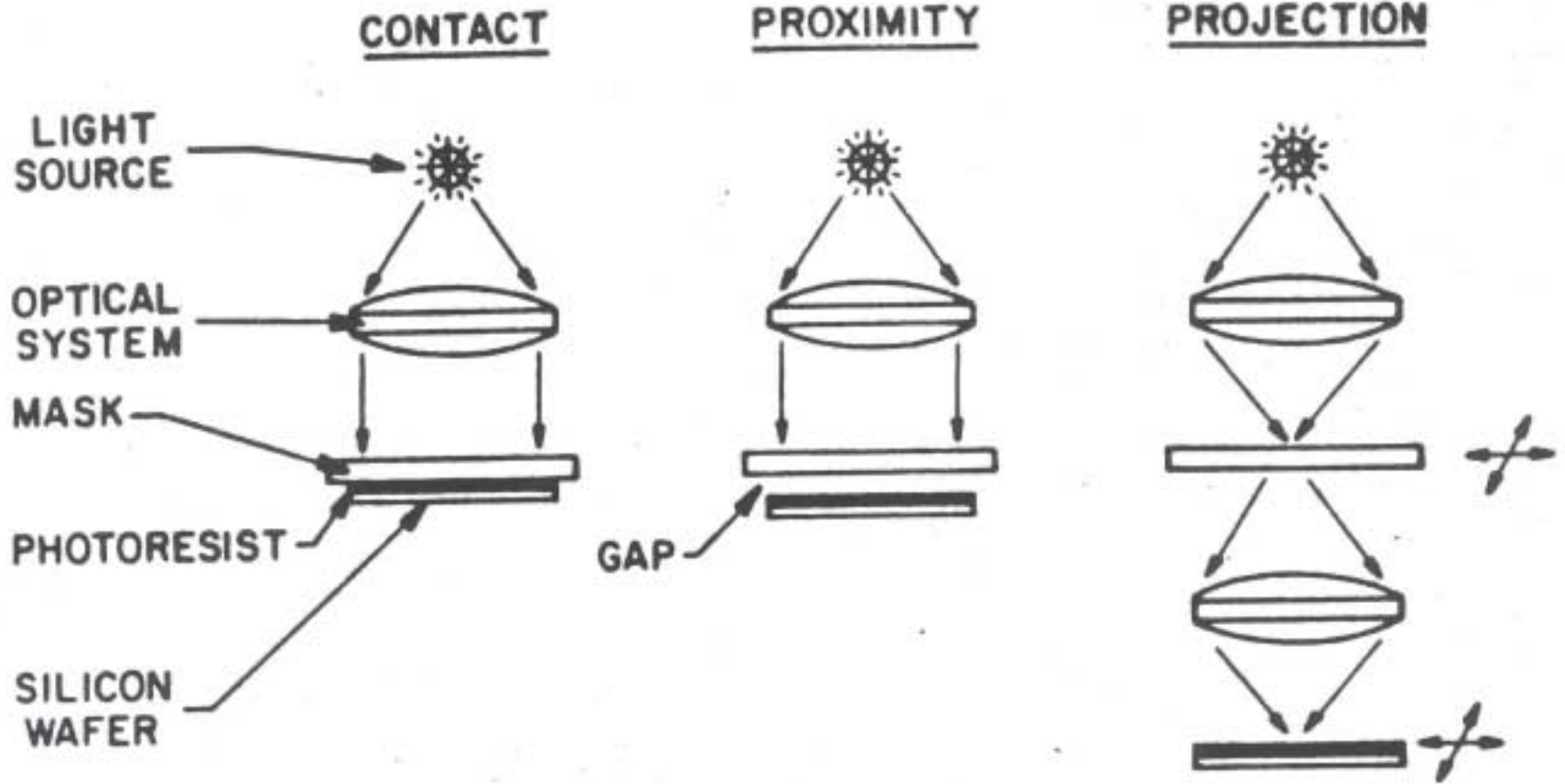
- Solvents: Most spinning solvents are based on esters of ethyleneglycol (Cellosolve, Cellosolve acetates). But these are carcinogens. There has been a switch to esters of linear high molecular weight acids.
- Criteria for choosing solvents:

- 1) Polymer must have a large radius of gyration (R_g) in the solvent. Radius of gyration is the mean square end to end distance. Larger the R_g , the more extended the polymer is and the better it is for spinning. If R_g is small, the polymer eventually becomes immiscible and precipitates.



- 2) Polymer must remain in solvent in the entire spinning process otherwise aggregation leads to pinholes and opaque spots. As solvent evaporates c and η increase. Solvent volatility must be low enough that c and η do not change rapidly during spinning.

Three methods of “printing”



Contact & Proximity printing

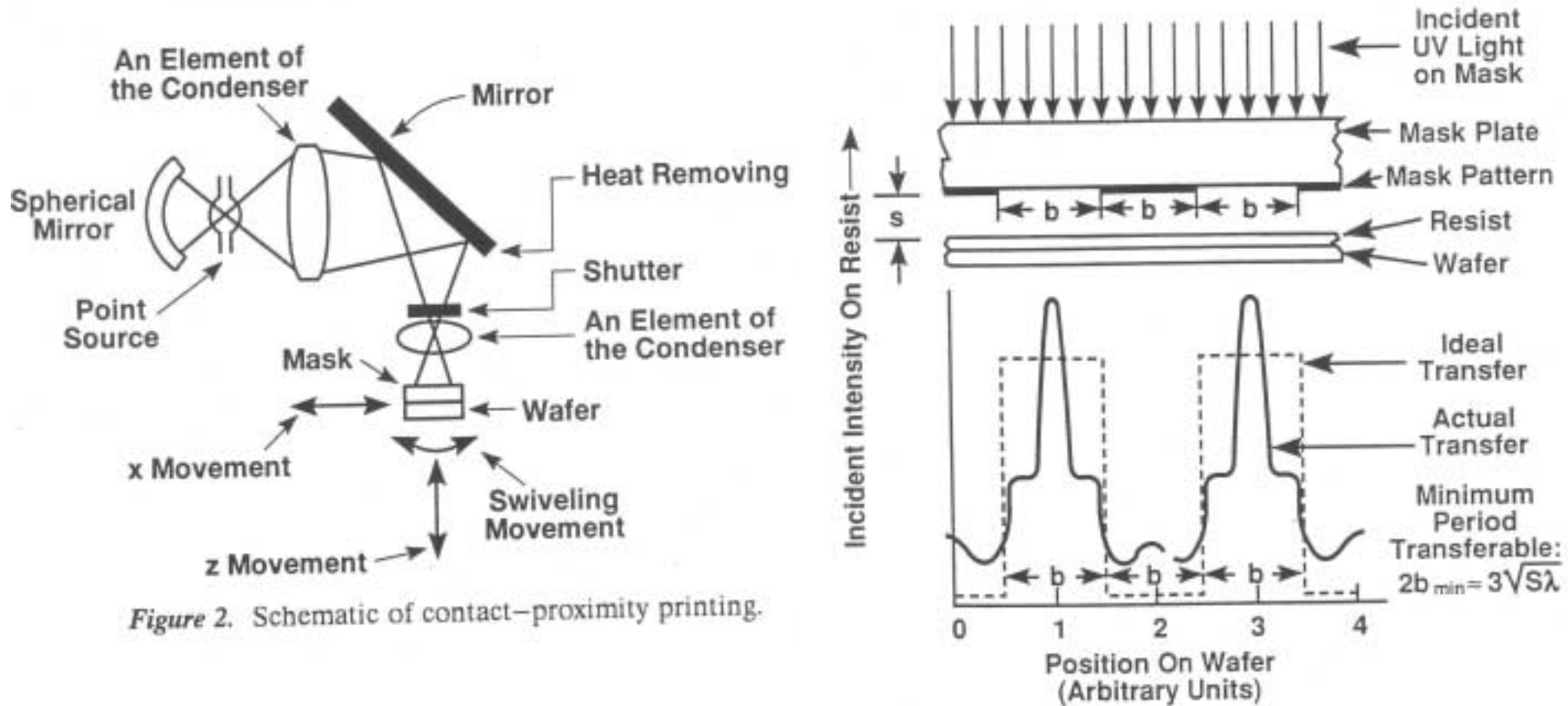
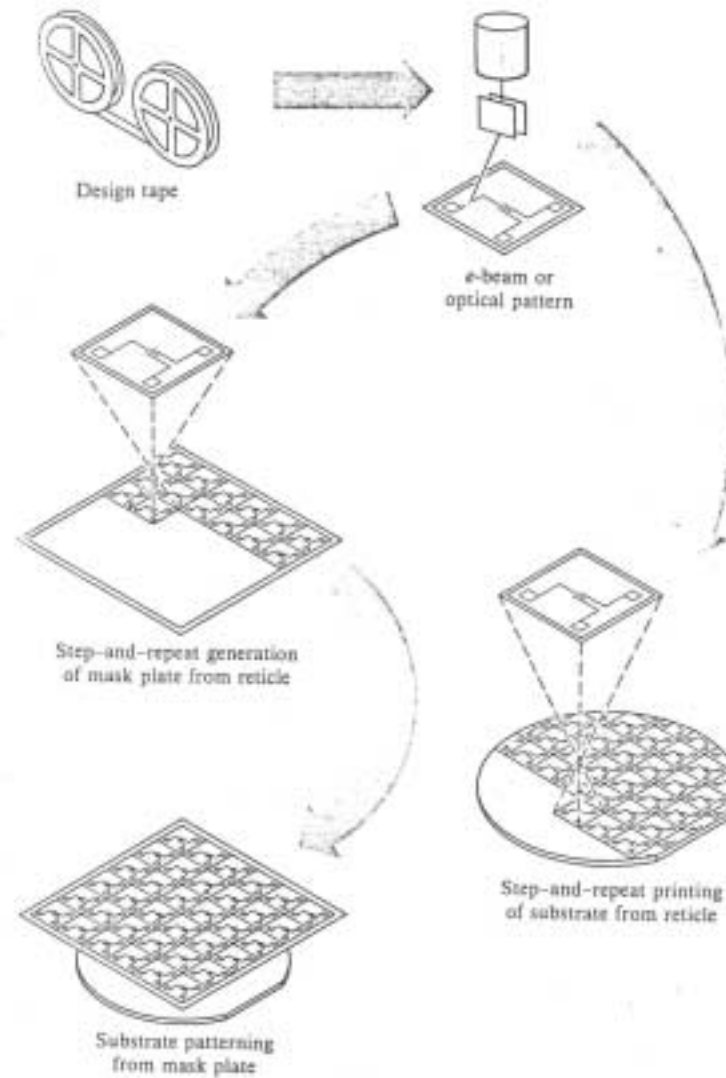
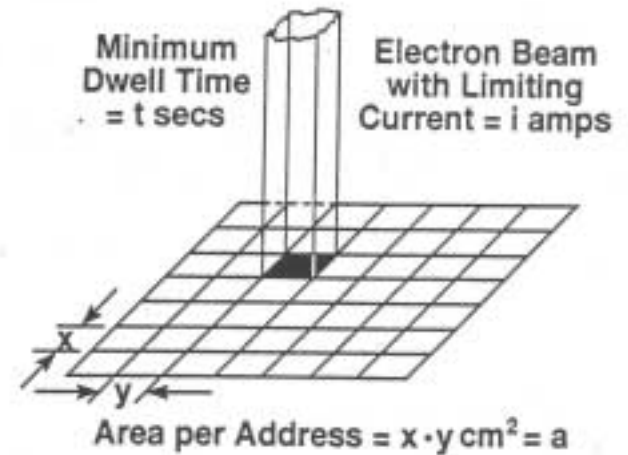
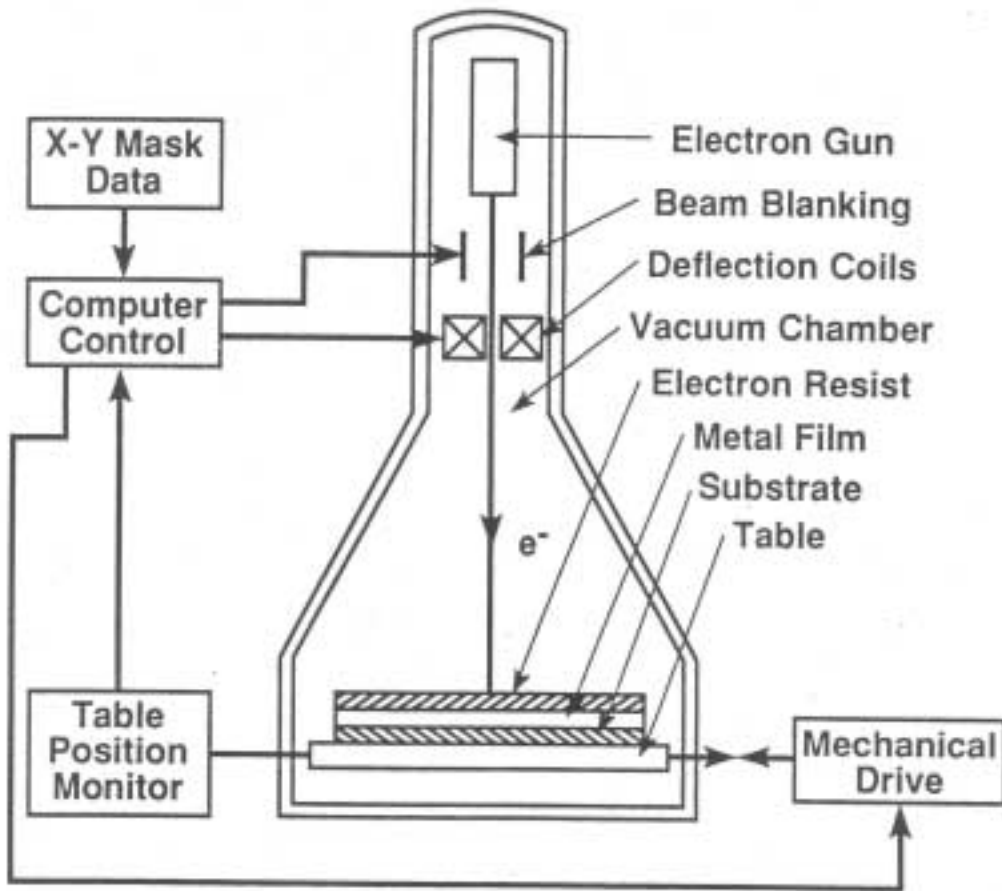


Figure 2. Schematic of contact-proximity printing.

Step Scan



E-beam lithography $E=h\nu=hc/\lambda$



Evolution of Lithographic methods

