

Micro- and Nano- Fabrication and Replication Techniques



Why do we have to write thing
small and replicate fast ?





Plenty of Room at the Bottom

Richard P. Feynman, December 1959

How do we *write* it? We have no standard technique to do this now. But let me argue that it is not as difficult as it first appears to be. We can **reverse the lenses of the electron microscope** in order to demagnify as well as magnify. A source of **ions**, sent through the microscope lenses in reverse, could be focused to a very small spot. We could write with that spot like we write in a TV cathode ray oscilloscope, by going across in lines, and having an adjustment which determines the amount of material which is going to be deposited as we scan in lines. **This method might be very slow** because of space charge limitations. There will be more rapid methods. We could first make, perhaps by some **photo process**, a screen which has holes in it in the form of the letters. Then we would strike an arc behind the holes and draw metallic ions through the holes; then we could again use our system of lenses and make a small image in the form of ions, which would deposit the metal on the pin.





Plenty of Room at the Bottom

Richard P. Feynman, December 1959

A simpler way might be this (though I am not sure it would work): **We take light and, through an optical microscope running backwards**, we focus it onto a very small photoelectric screen. Then electrons come away from the screen where the light is shining. These electrons are focused down in size by the electron microscope lenses to impinge directly upon the surface of the metal. Will such a beam etch away the metal if it is run long enough? I don't know. If it doesn't work for a metal surface, it must be possible to find some surface with which to coat the original pin so that, where the electrons bombard, a change is made which we could recognize later.





Plenty of Room at the Bottom

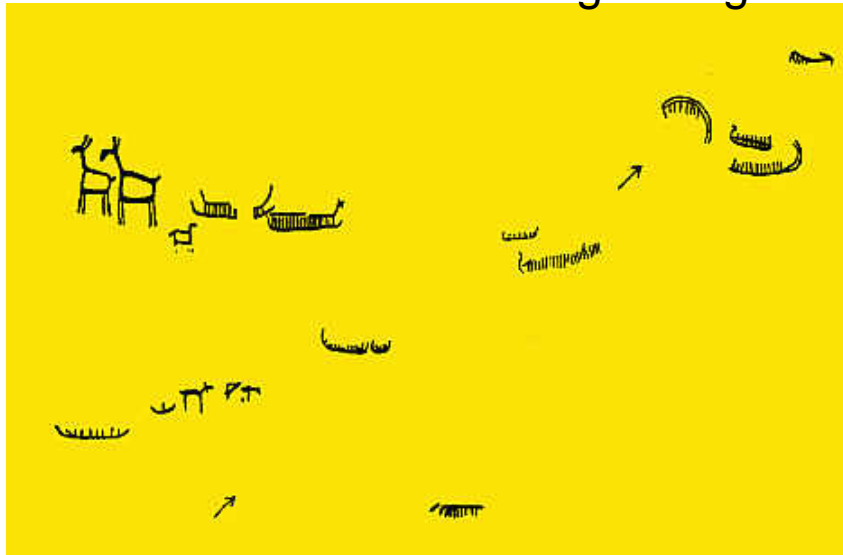
Richard P. Feynman, December 1959

There is no intensity problem in these devices---not what you are used to in magnification, where you have to take a few electrons and spread them over a bigger and bigger screen; it is just the opposite. The light which we get from a page is concentrated onto a very small area so it is very intense. The few electrons which come from the photoelectric screen are demagnified down to a very tiny area so that, again, they are very intense. **I don't know why this hasn't been done yet!**



Ancient Patterning

"This is the Elks' land". A greeting at the mouth of Dalbergsaa, Southern Dal.



It seems that the carvings of Northern Scandinavia's including Kola Peninsula are the oldest.

Large figures of ritual animals characterise the Mesolithic period mostly before c.4200 BC



Writing by Inks



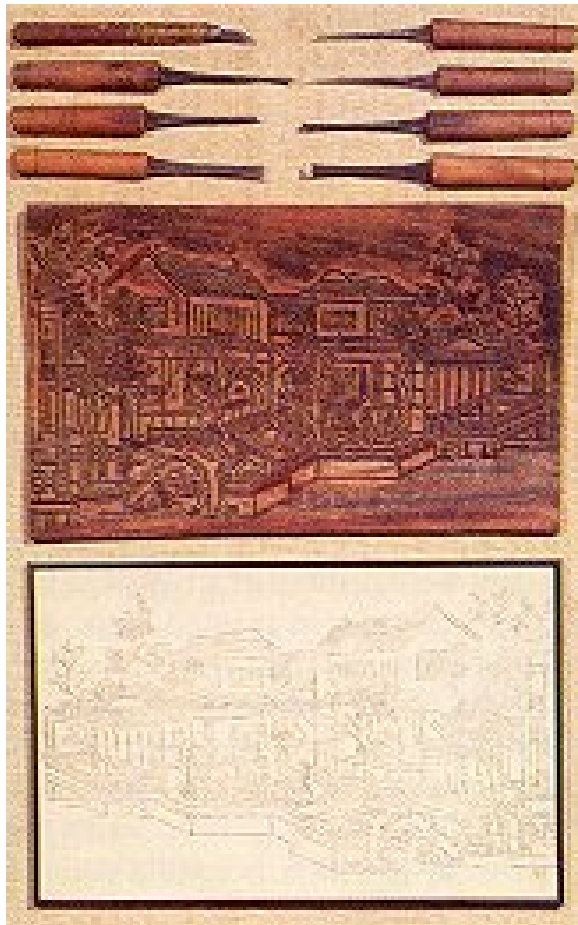
Writing Brush ~2000 Ys



Quill Pen ~1000 Ys



Chinese Replication



868



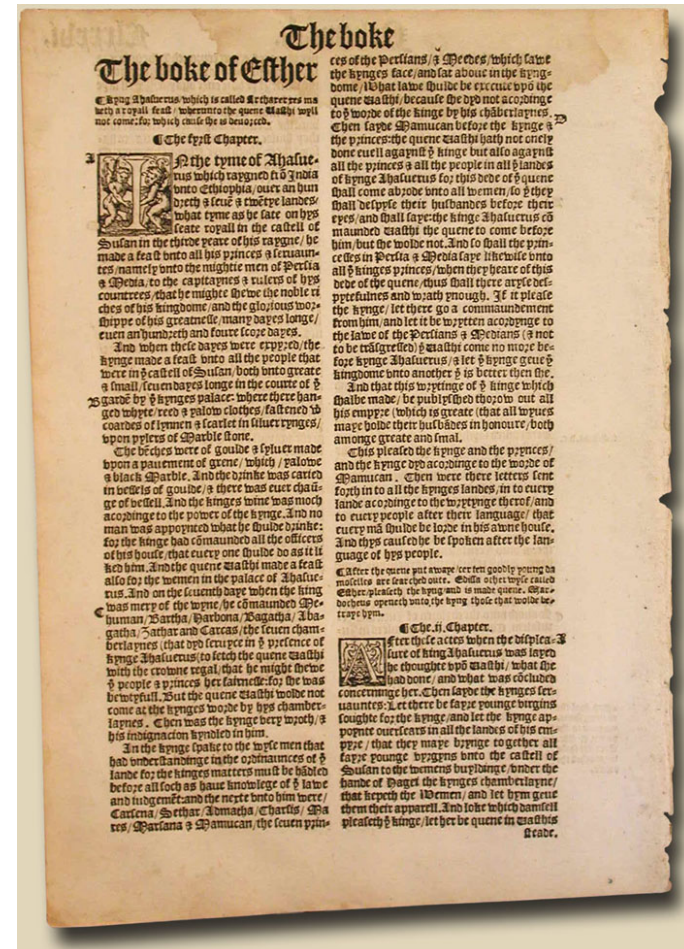
1041 and 1048



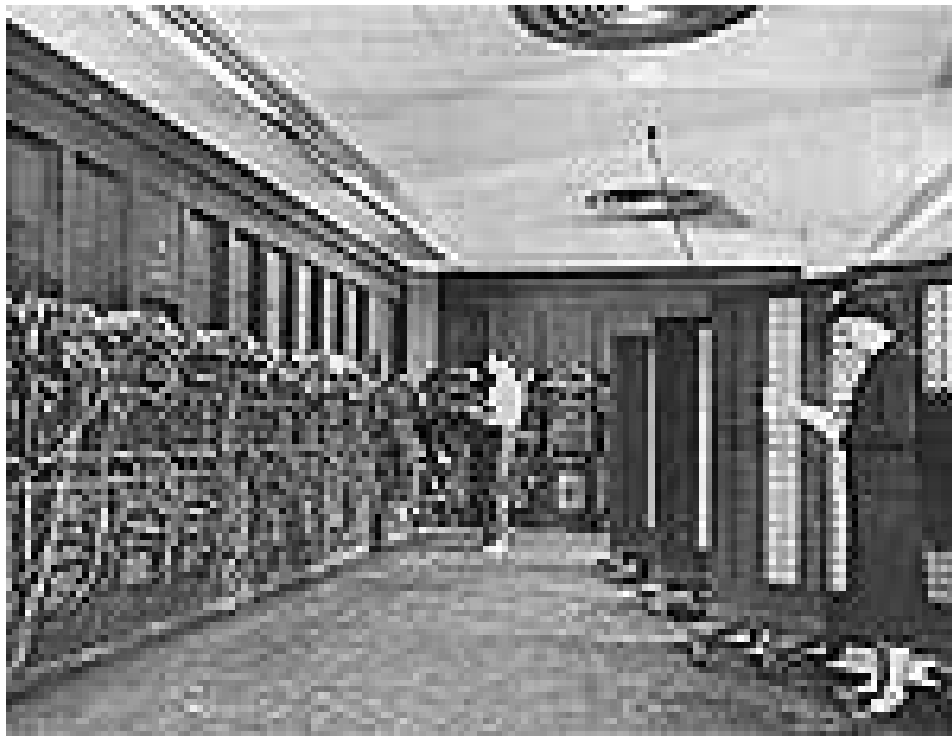
Western Replication



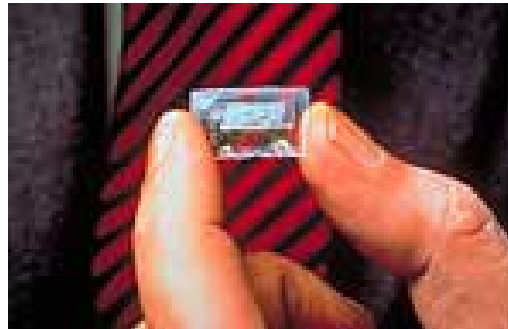
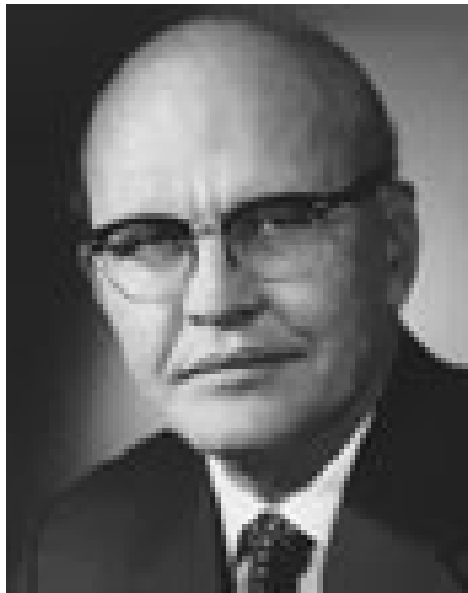
1543 - 1610



Building a computer



First Integrated Circuit

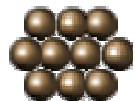
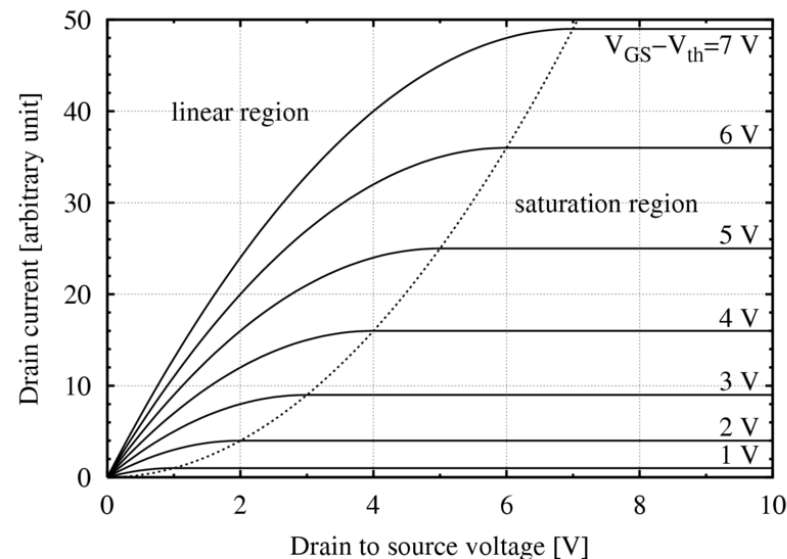
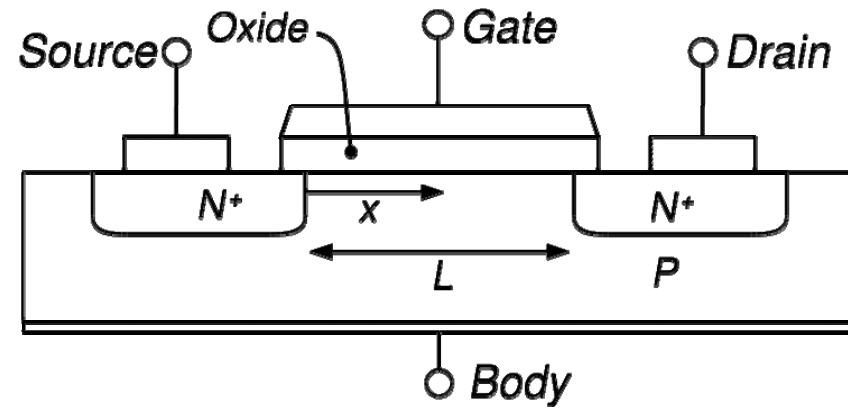
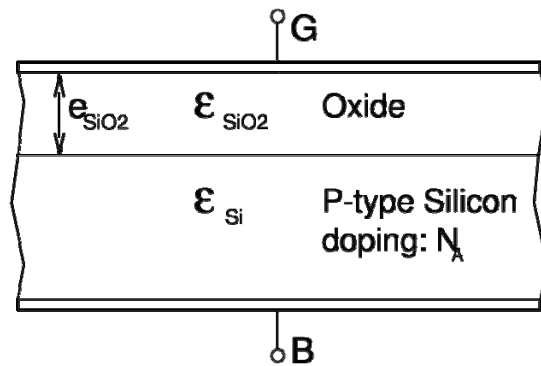


"What we didn't realize then was that the integrated circuit would reduce the cost of electronic functions by a factor of a million to one, nothing had ever done that for anything before" - Jack Kilby 2000 Nobel Prize

1958 Texas Instruments



metal-oxide-semiconductor field-effect transistor (MOSFET)



Fabrication Techniques

- Laser writing
- E-beam writing
- Focus ion beam writing
- Nonlinear optical writing
- Interference lithography
- Dip-pen lithography
- SPM lithography

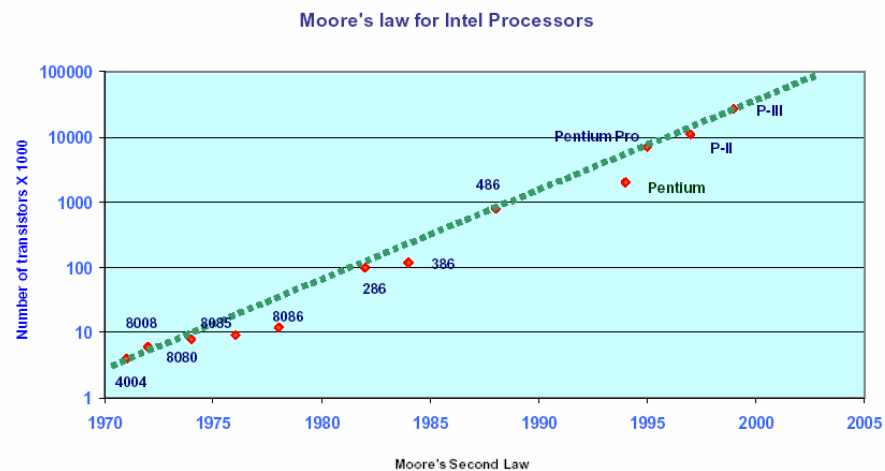


Replication Techniques

- Photolithography
- X-ray or e-beam projection
- Microcontact printing (soft-lithography)
- Nanoimprint



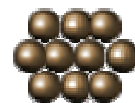
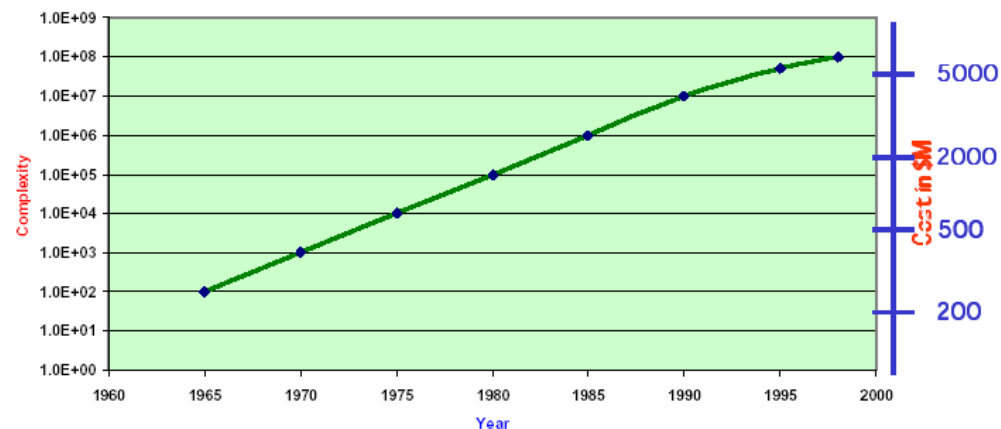
Moore's Law



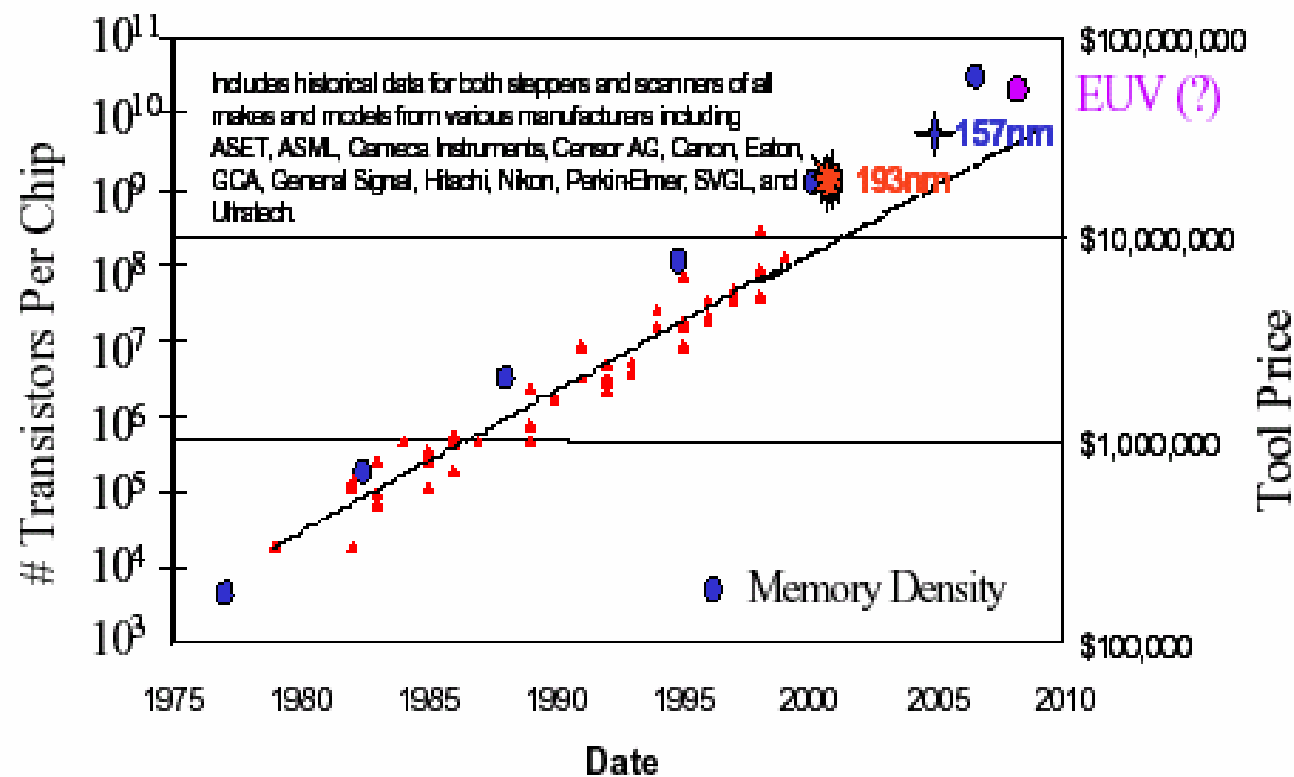
Moore's Laws

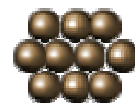
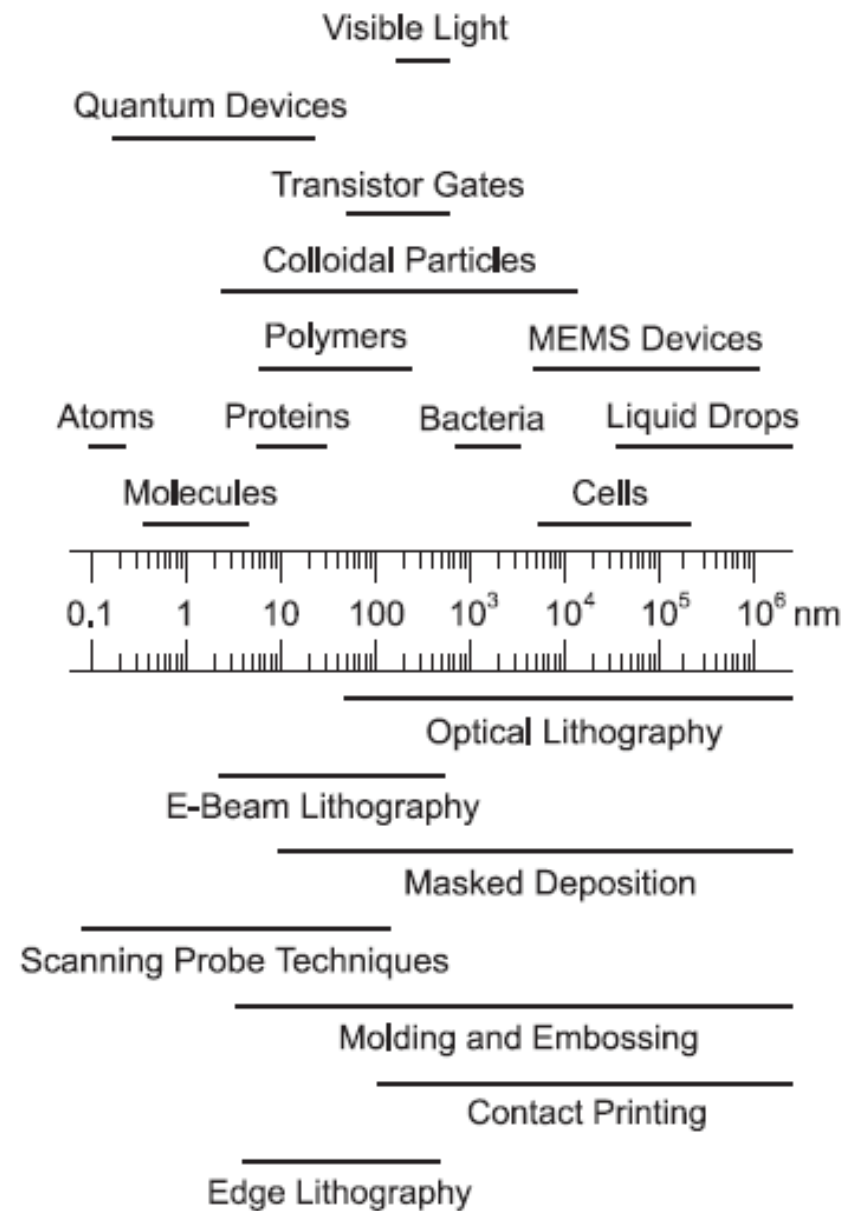
First Law: Number of components in a chip (IC) will double roughly every 18 months (1965, in *Electronics*). This has held true more or less since then.

Second Law: Facility costs increase on a semilog scale (terminology due to Eugene Meieran, Intel Fellow). Fab costs double approximately every four years.

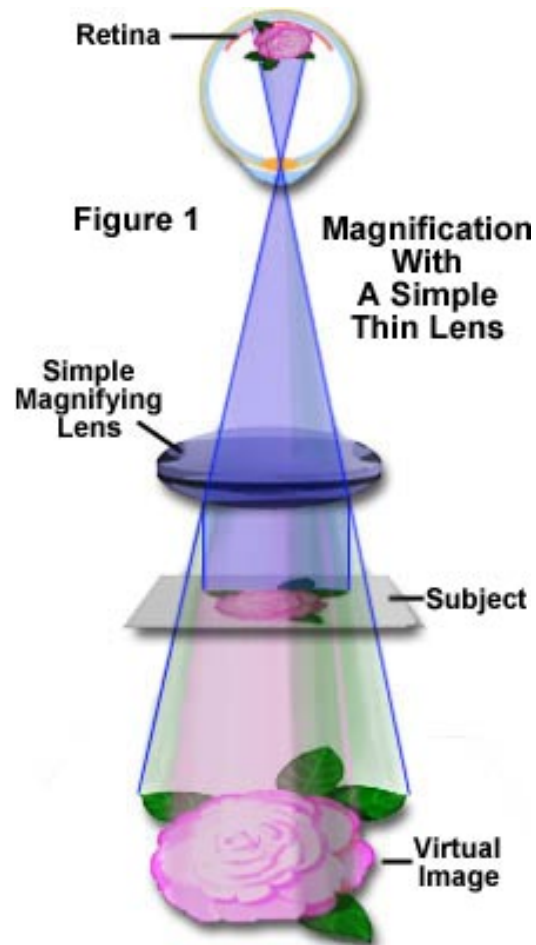


Tool Cost

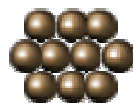
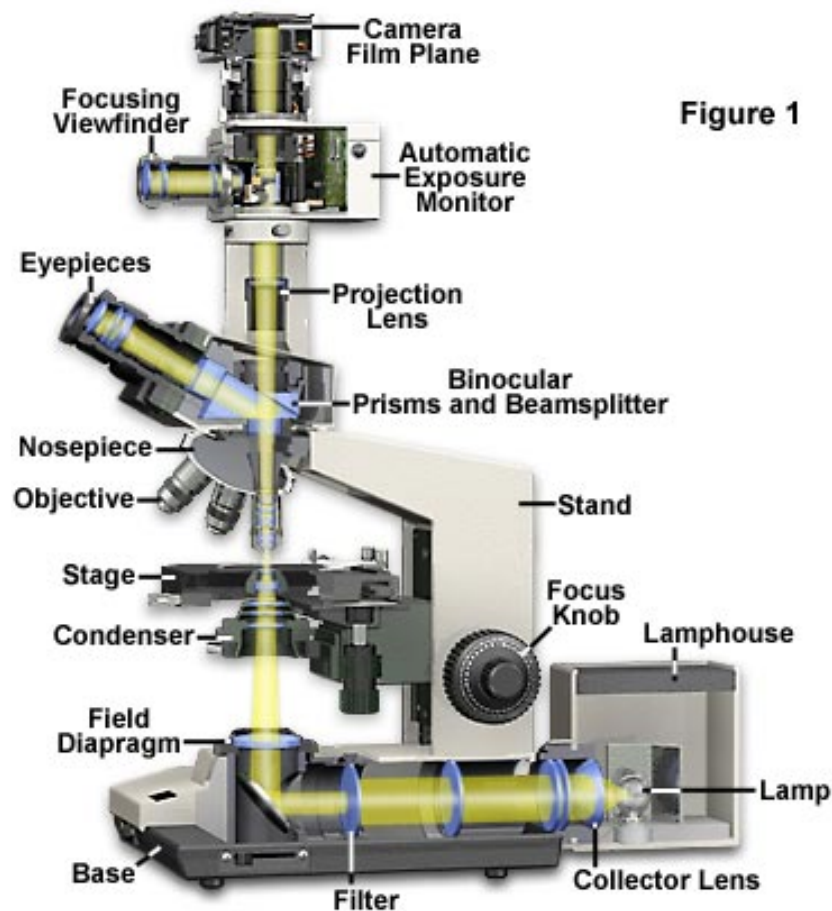




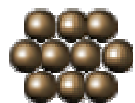
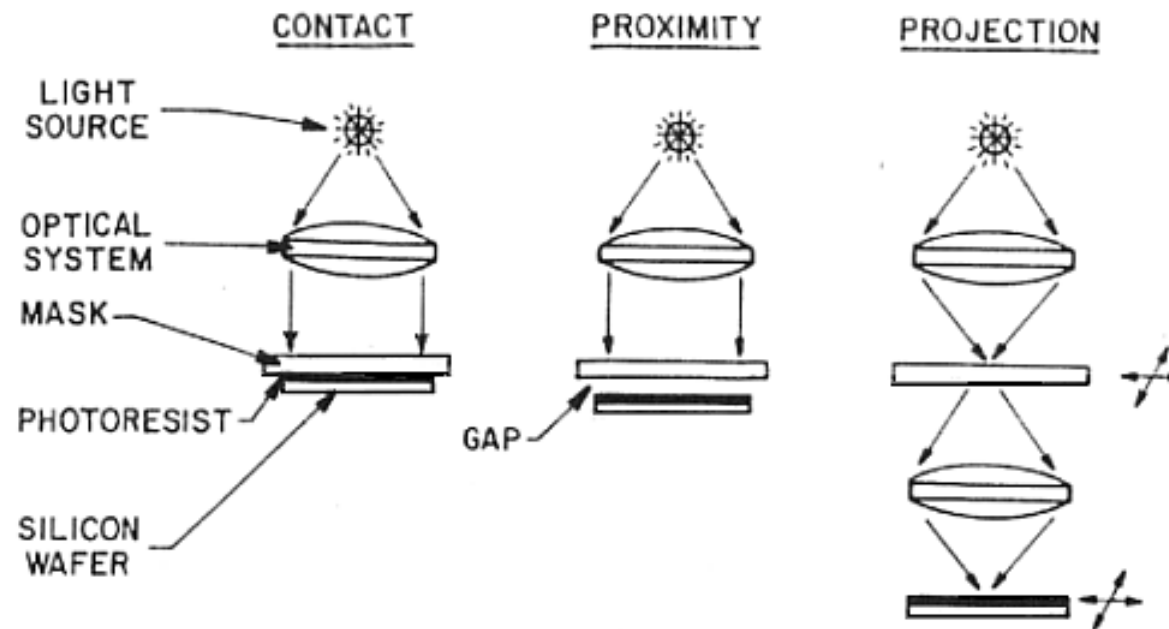
Optical Microscope



Modern Microscope Component Configuration

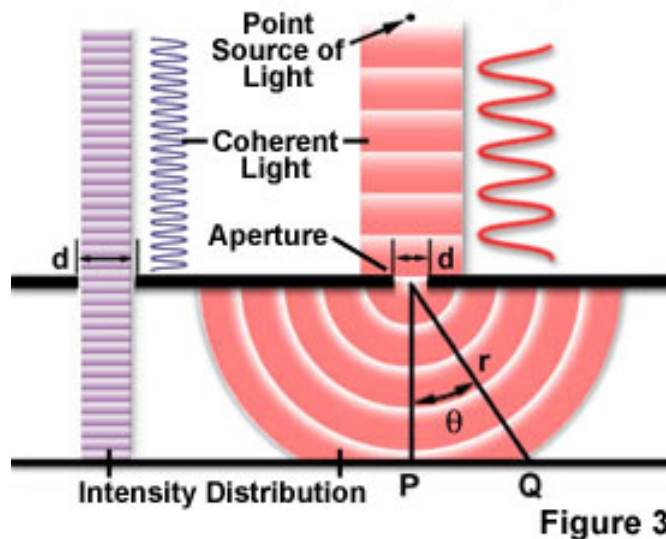


Methods of Photolithography

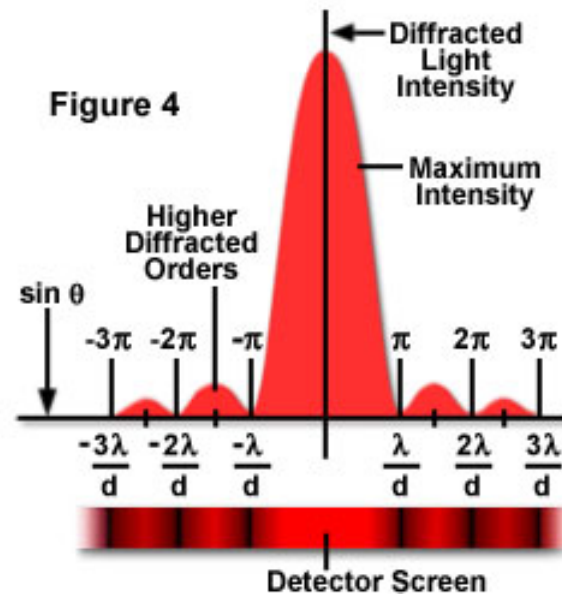


Limit of Photolithography

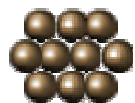
Diffraction of Light Through an Aperture



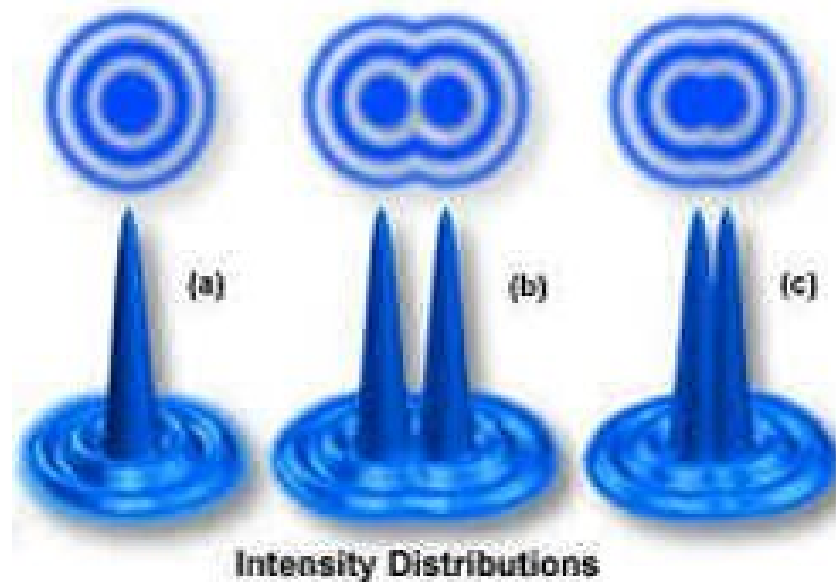
Intensity Distribution of Diffracted Light



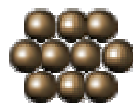
$$r = 1.22 \times \lambda / (2 \times \text{N.A.})$$
$$\text{N.A.} = n \times \sin(\theta)$$



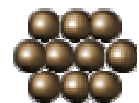
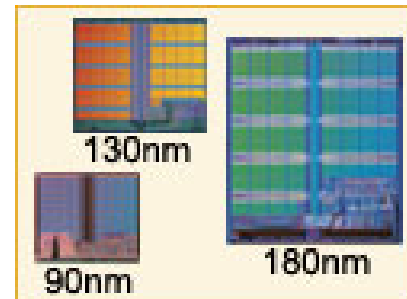
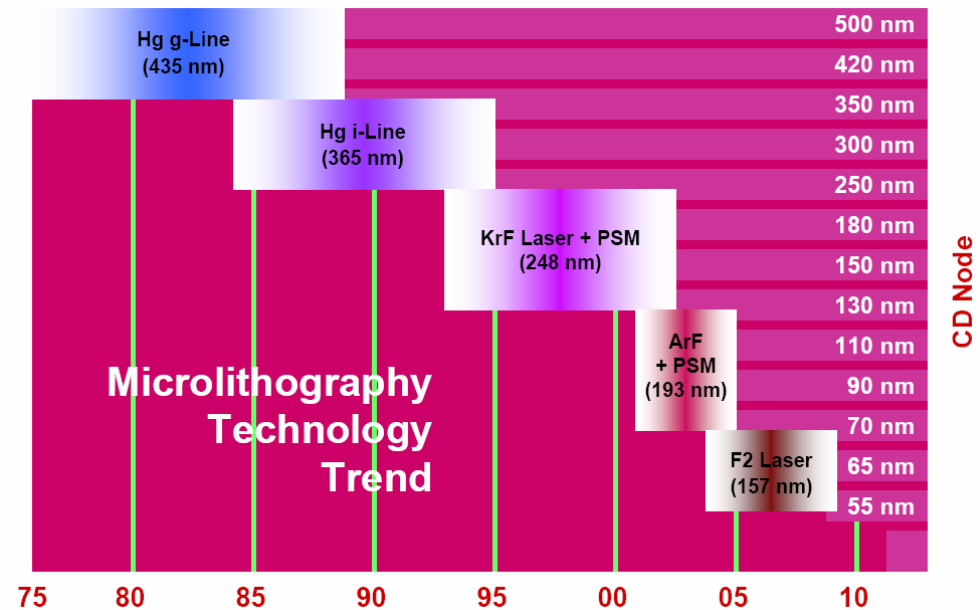
Diffraction Limit



$$\text{Resolution} = K \times \lambda / (\text{N.A.})$$
$$\text{Depth of Focus} = \lambda / (\text{N.A.})^2$$
$$K = 0.61$$



Photolithography



UV Wavelength (nm)	Wavelength Name	UV Emission Source
436	g-line	Mercury arc lamp
405	h-line	Mercury arc lamp
365	i-line	Mercury arc lamp
248	Deep UV (DUV)	Mercury arc lamp or Krypton Fluoride (KrF) excimer laser
193	Deep UV (DUV)	Argon Fluoride (ArF) excimer laser
157	Vacuum UV (VUV)	Fluorine (F ₂) excimer laser

Year	Linewidth (nm)	Wavelength (nm)
1986	1,200	436
1988	800	436/365
1991	500	365
1994	350	365/248
1997	250	248
1999	180	248
2001	130	248
2003	90	248/193
2005 (fcst)	65	193
2007 (fcst)	45	193



Water Immersion Lithography

Year	Linewidth (nm)	Wavelength (nm)
1986	1200	436 g-line mercury lamp
1988	800	436/365
1991	500	365 i-line mercury lamp
1994	350	365/248
1997	250	248 KrF excimer laser
1999	180	248
2001	130	248
2003	90	248/193
2005	65	193 ArF excimer laser
2007	45	193/157

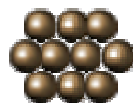
Resolution (R) = $K \times \lambda / (N.A.)$

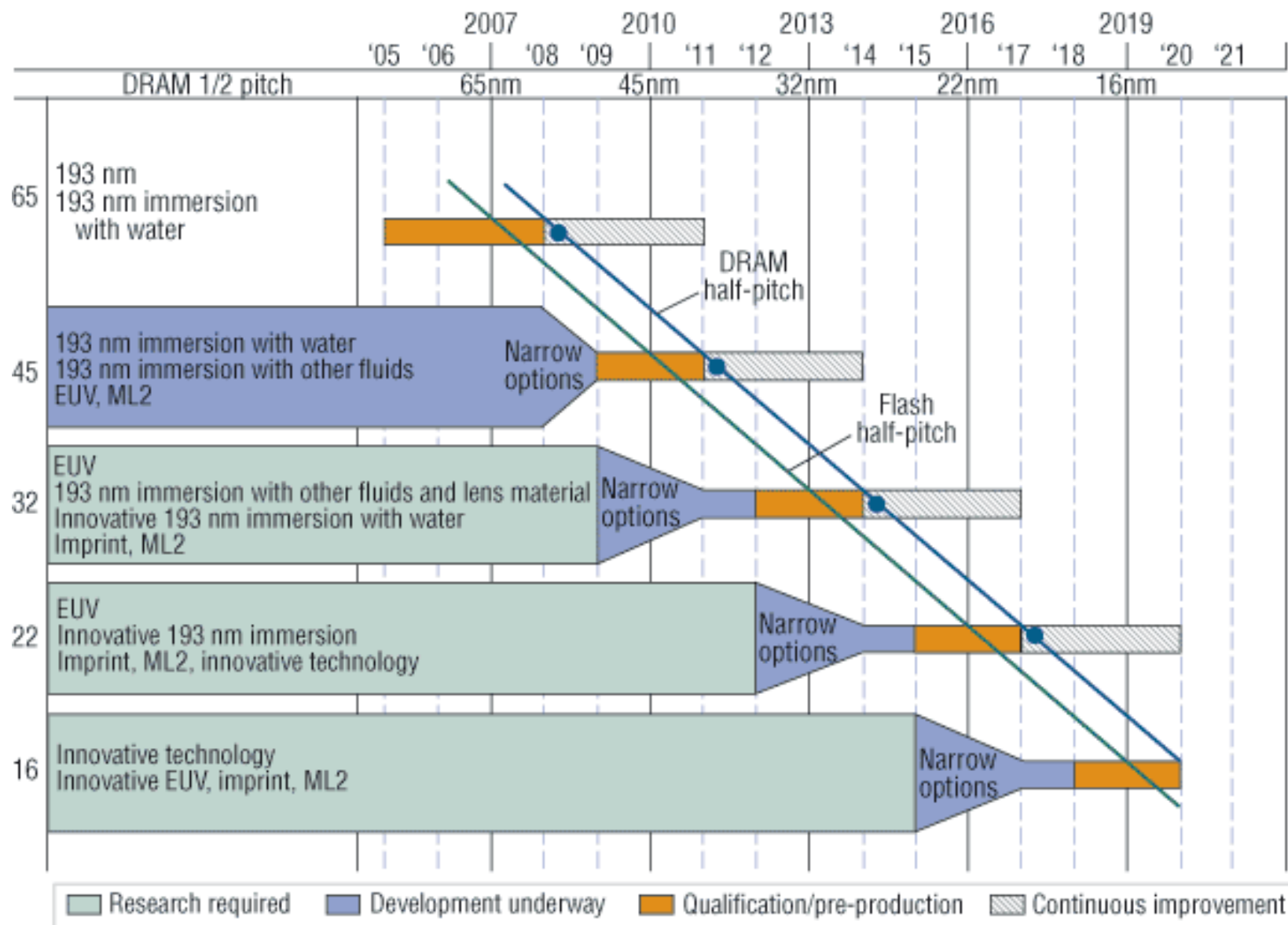
$K = 0.25$, $NA \sim 1.4$, $\lambda = 193$

$R = 35 \text{ nm}$

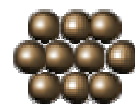
Air $n = 1.0003$

Water $n = 1.437$

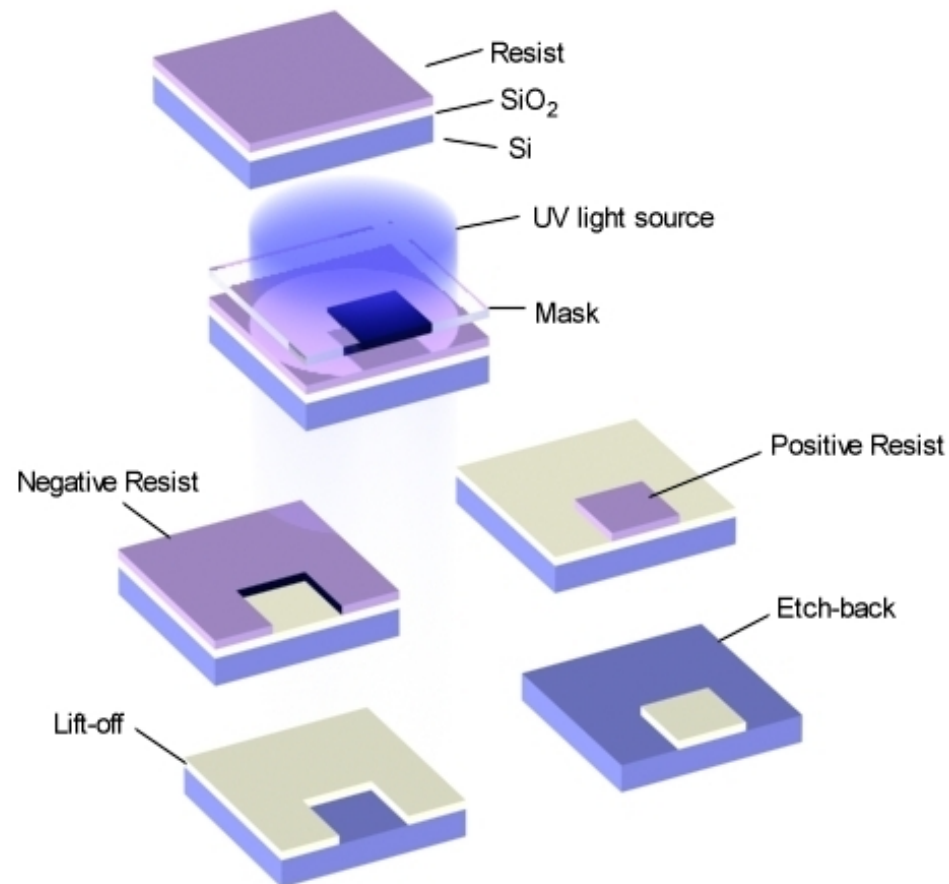




This legend indicates the time during which research, development, and qualification/pre-production should be taking place for the solution.

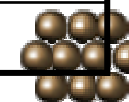


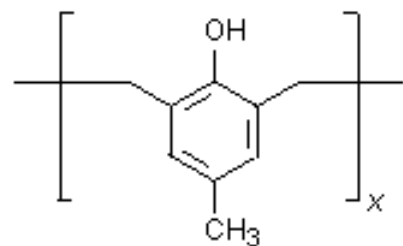
Photolithography Process



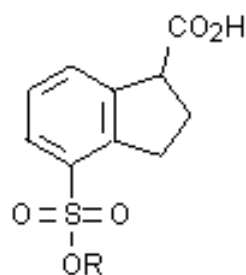
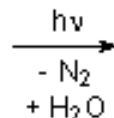
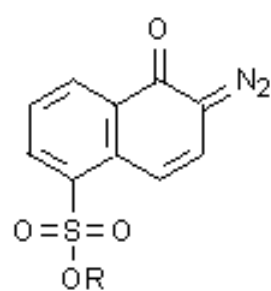
RCA Cleaning (By Radio Corporation of America in 1965)

Chemicals	Volume ratio	Procedure Time (min)	Operation Temperature	Function
Trichloroethane		5	Room T	Dissolve Organic
Acetone		5	Room T	Dissolve Organic
DI Water		5	Room T	Washing
H ₂ SO ₄ (98%)-H ₂ O ₂ (30%) (Piranha Solution)	3:1	10-20	~90°C	Oxide and Dissolve Organic and Metals
DI Water		5	Room T	Washing
HF(49 wt %)-H ₂ O	~2:100	10-20	Room T	Dissolve surface SiO ₂
NH ₄ OH(29%)-H ₂ O ₂ (30%)- H ₂ O	1:1:5	10-20	~90°C	Oxide and Dissolve Metals
DI Water		5	Room T	Washing
HCl(37%)- H ₂ O ₂ (30%)- H ₂ O	1:1:5	10-20	~90°C	Oxide and Dissolve Metals
DI Water		5	Room T	Washing
Spin Dry (In lad – N ₂ blow)				



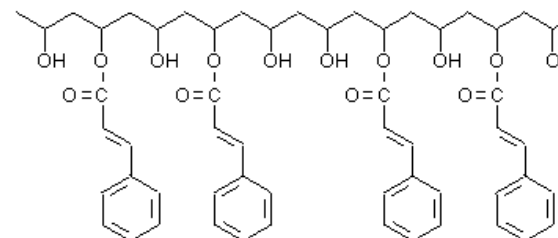
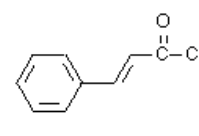
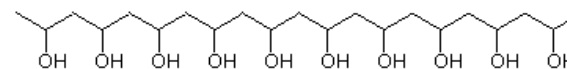
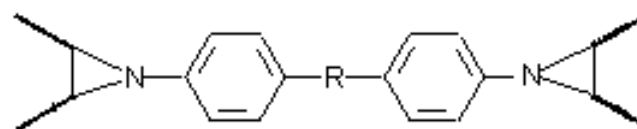
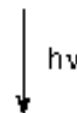
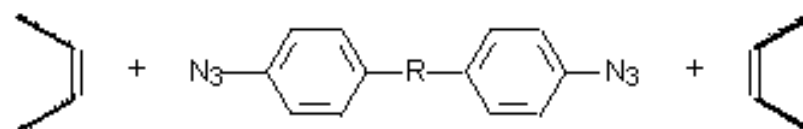


Novolak



Diazonaphthaquinone additive

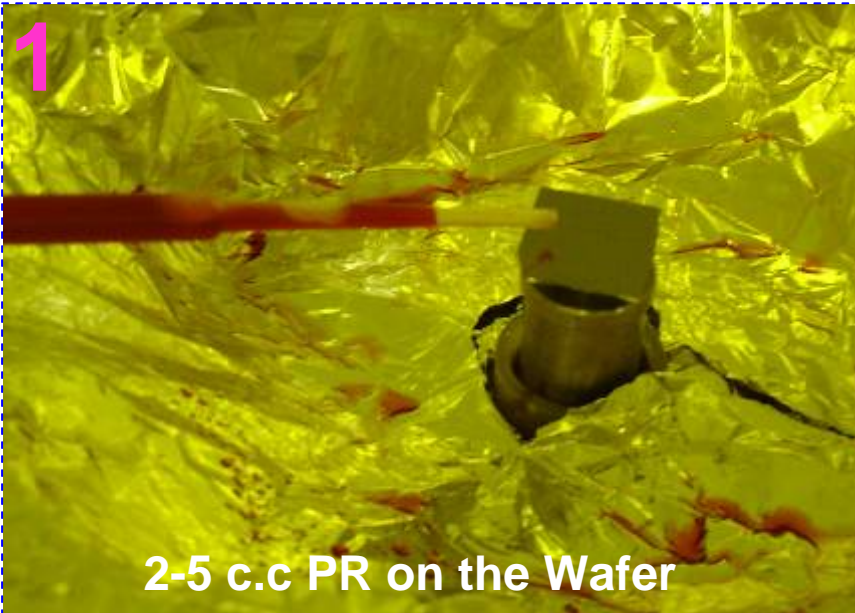
Positive tone



Negative tone

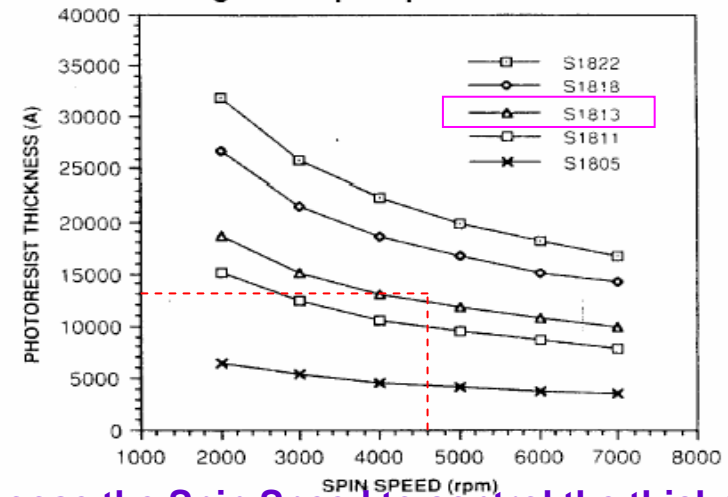


Spin Coating Photoresist on Wafer



3

MICROPOSIT S1800 PHOTO RESIST UNDYED SERIES
Figure 1. Spin Speed Curves



Choose the Spin Speed to control the thickness

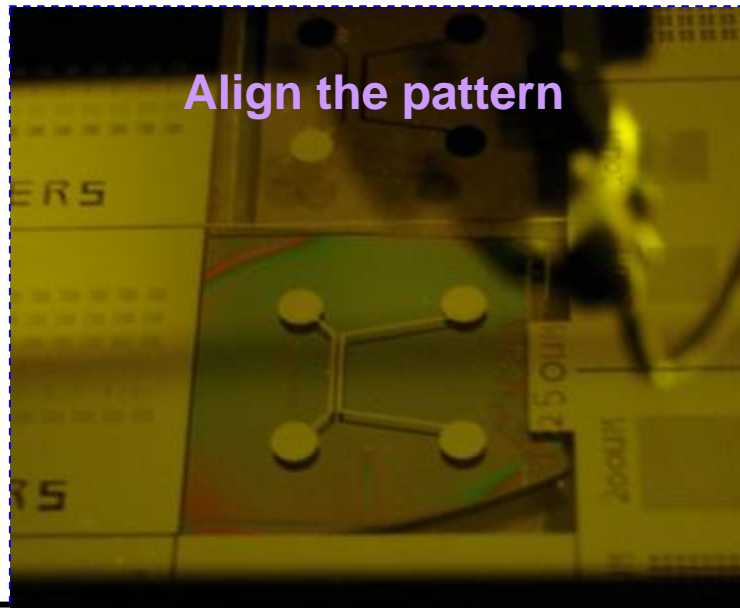
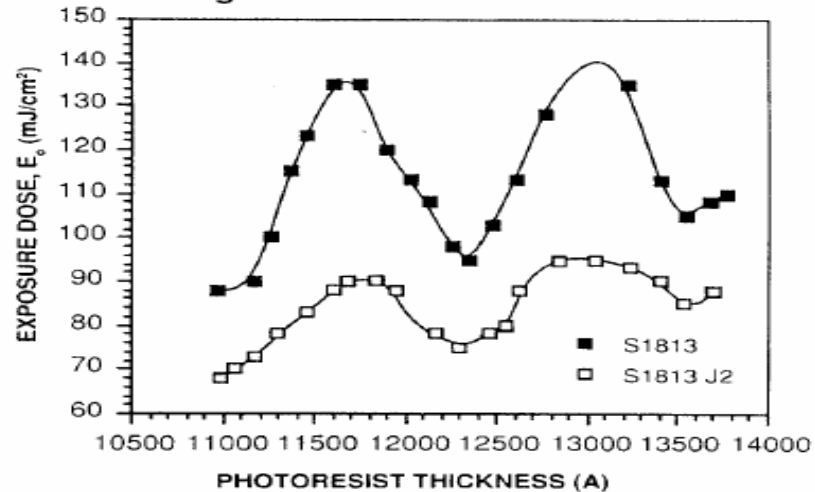
- Photo resist : Shipley 1813
- Spin : 2000 rpm 5 s
4500 rpm 15 s
- Soft Bake : 110°C 60 s
- Exposure : 7 s
- Developed : MF319 for 15 s

Standard Mask Size: 5" × 5"

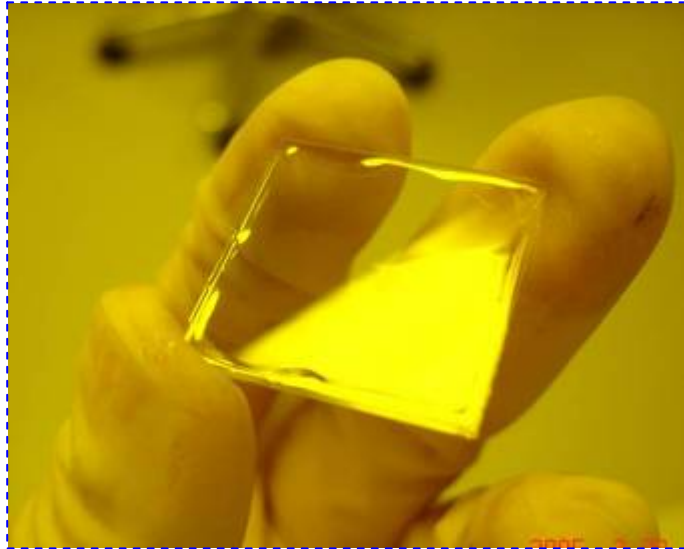


Align the pattern and Exposure

MICROPOSIT S1813 and S1813 J2 PHOTO RESISTS
Figure 4. Interference Curves



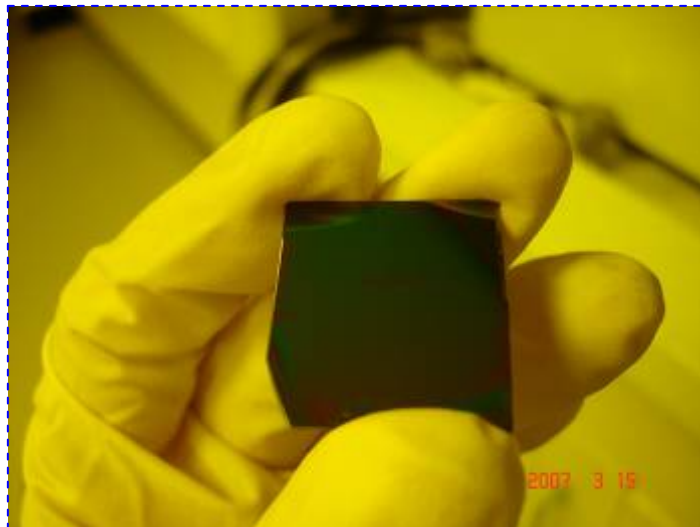
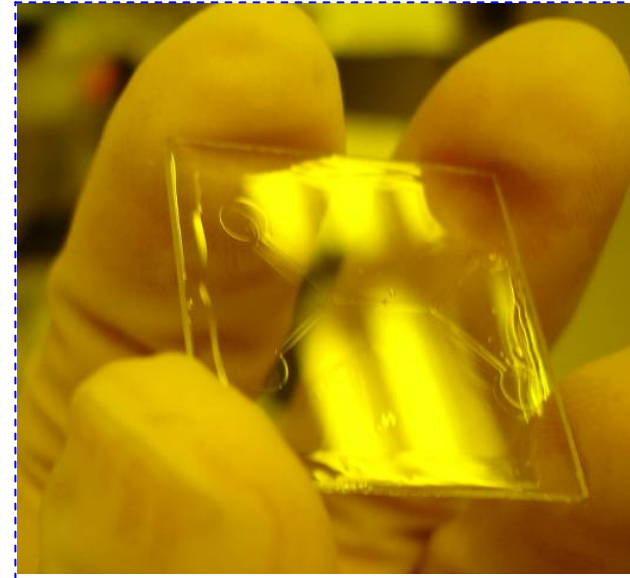
Some Photoresist Need PEB (post exposure Bake)



SU-8 2015

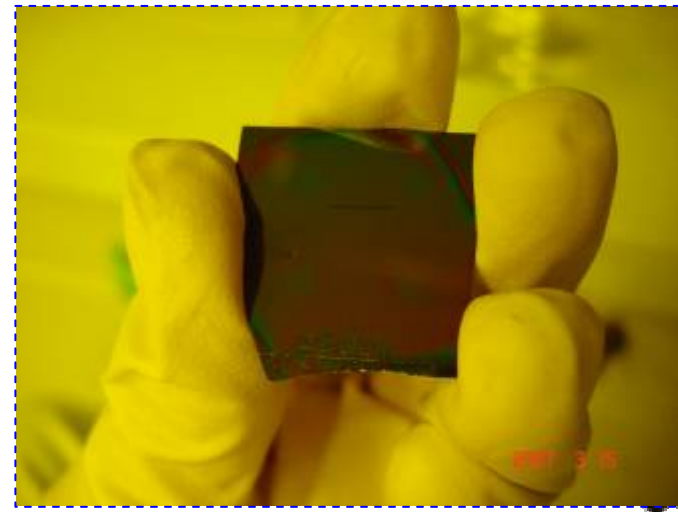
65 °C, 60 s

95 °C, 60 s

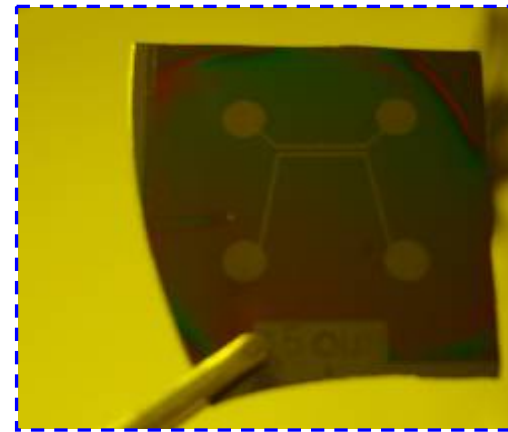
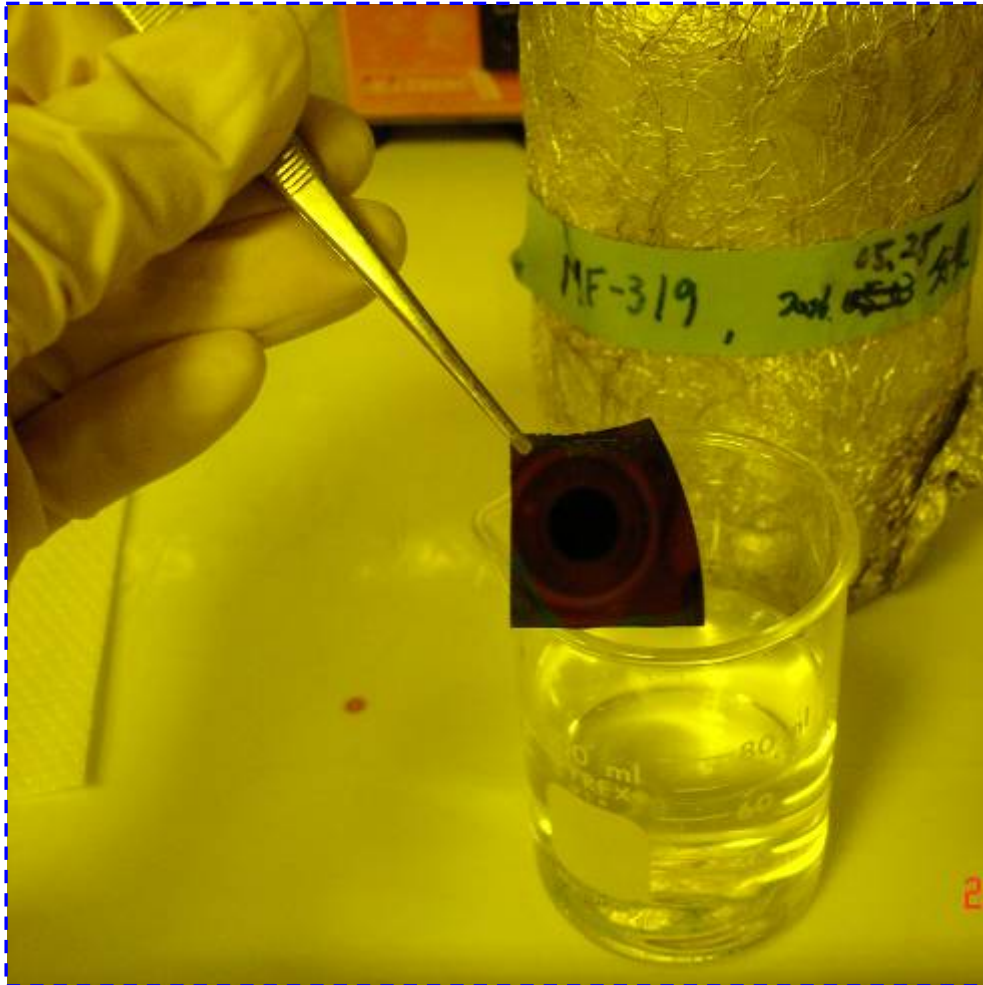


SPR 510A

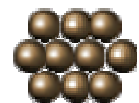
90 °C, 90 s



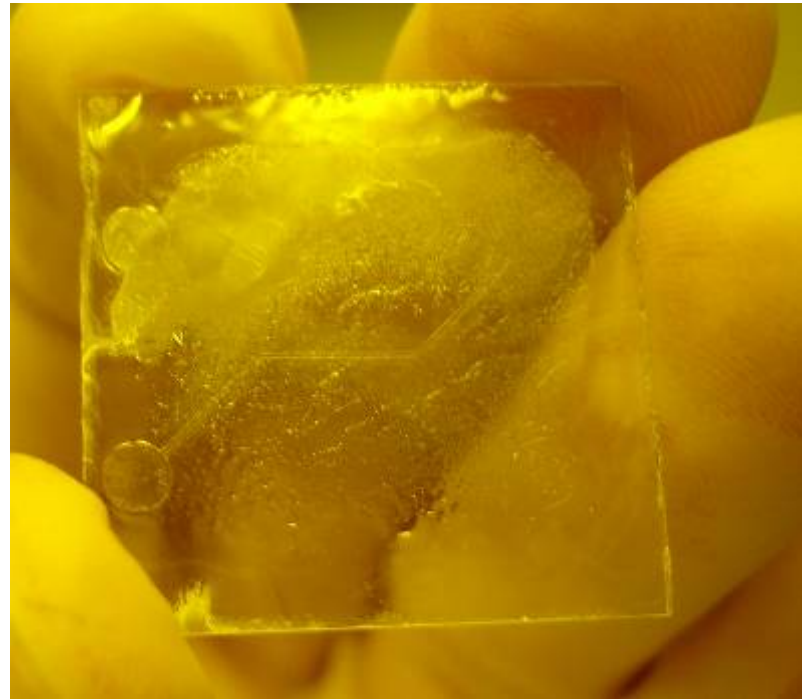
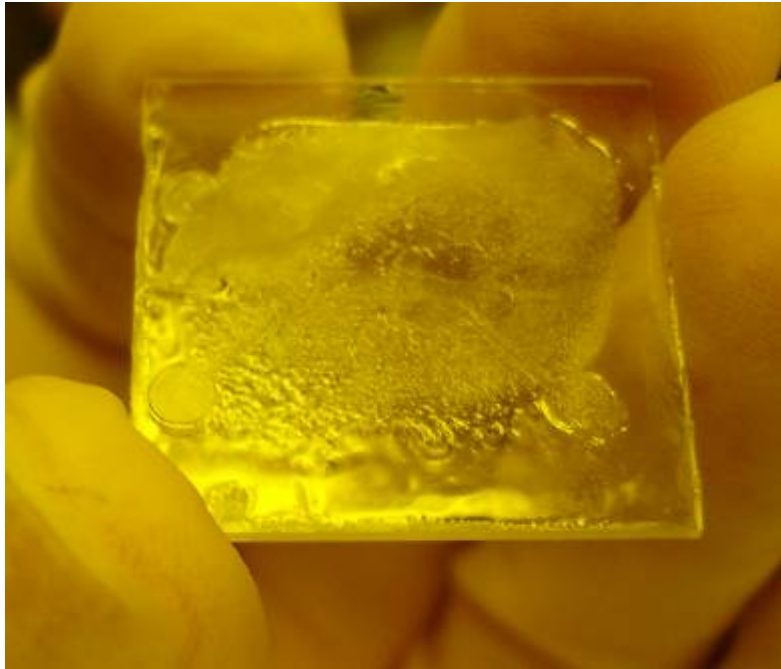
Develop the Photoresist



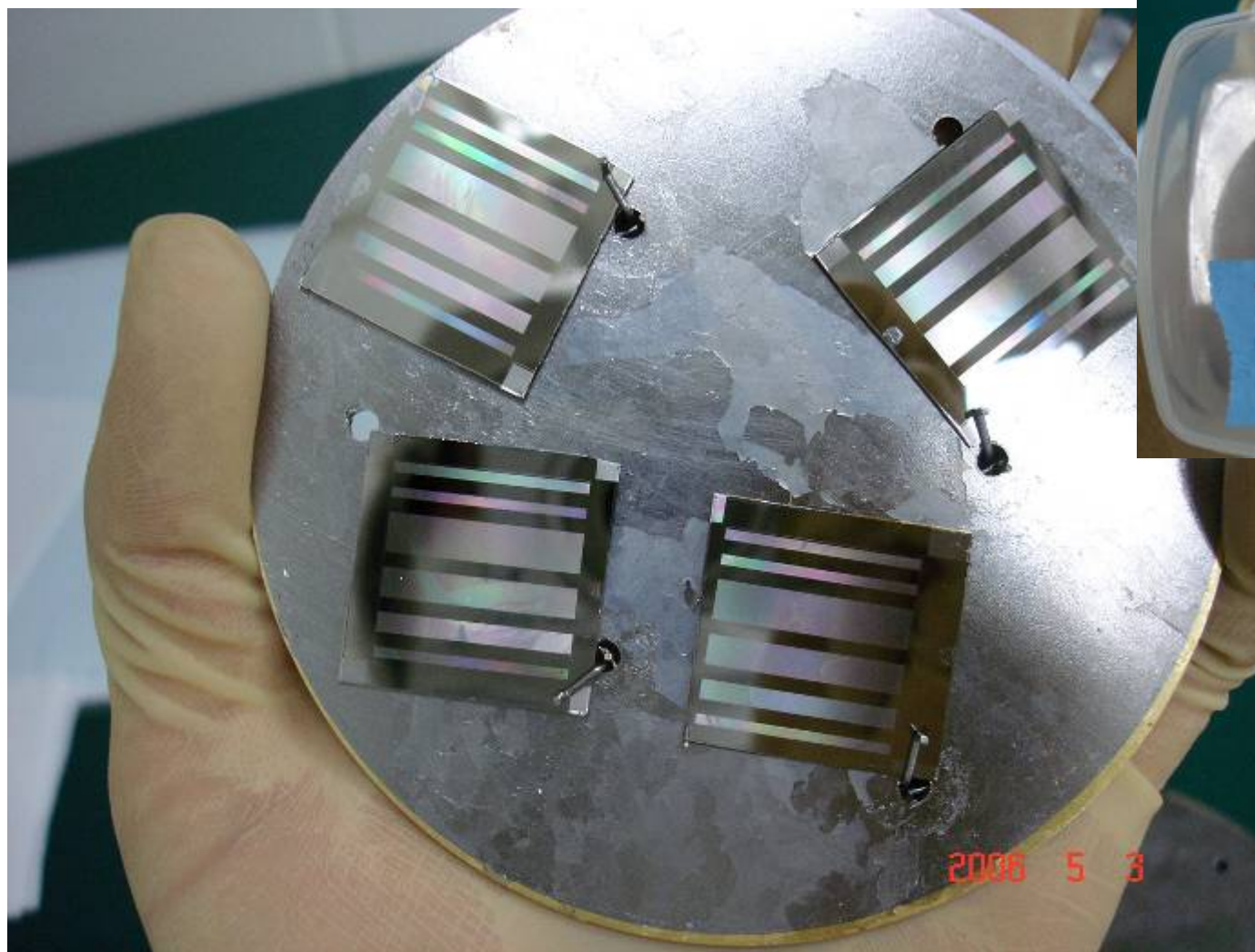
- Photo resist : **Shipley 1813**
- Spin : 2000 rpm 5 s
4500 rpm 15 s
- Soft Bake : 110°C 60 s
- Exposure : 7 s
- Developed : **MF319 for 15 s**



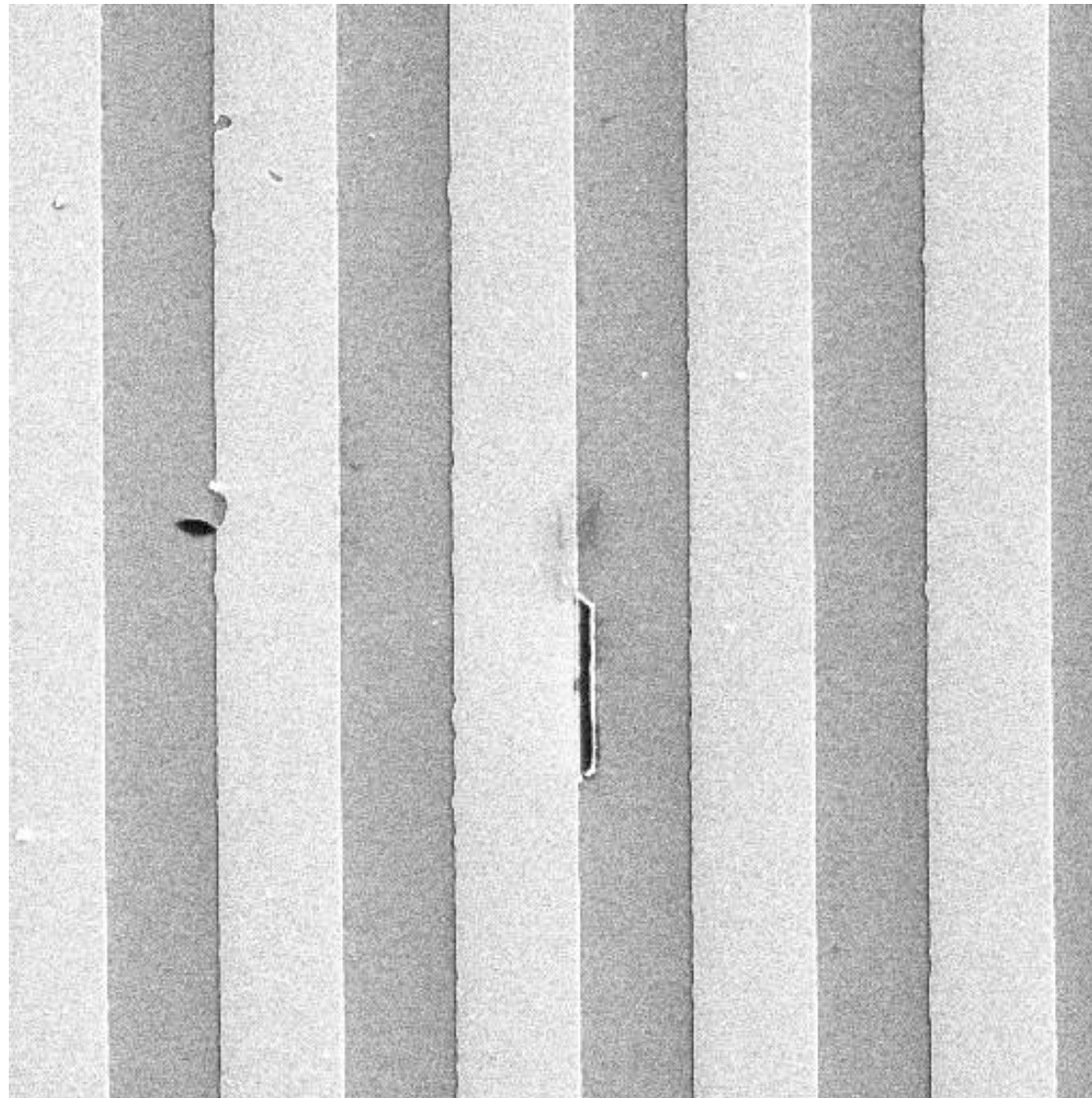
黃光室的黃光是很重要的



TI 50 nm

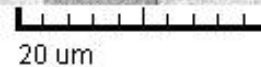


SEM image of Ti 50 nm on Si wafer

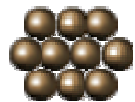


SEM MAG: 1.68 kx
View field: 89.76 μm
VAC: HiVac

DET: SE Detector
DATE: 05/04/06
SM: RESOLUTION



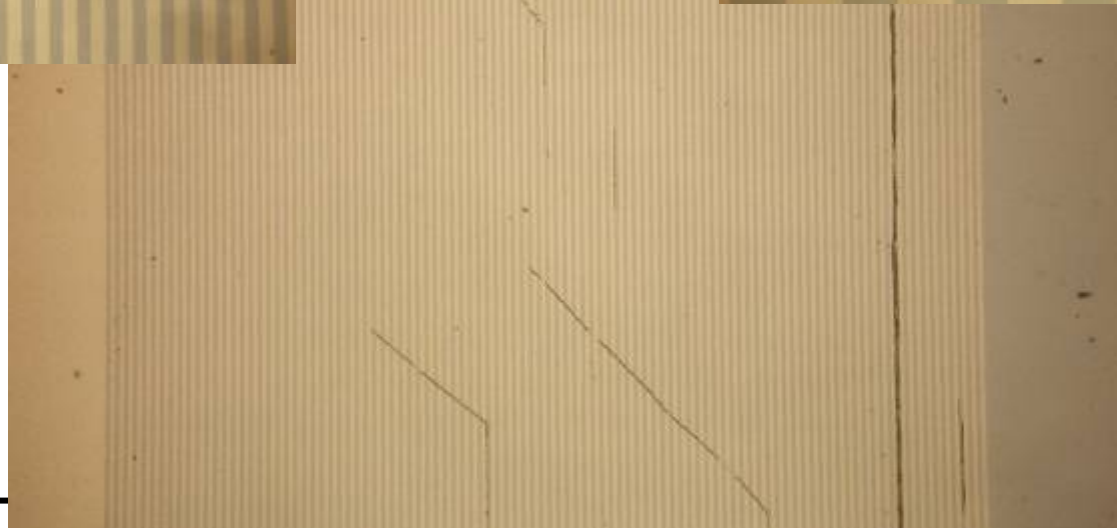
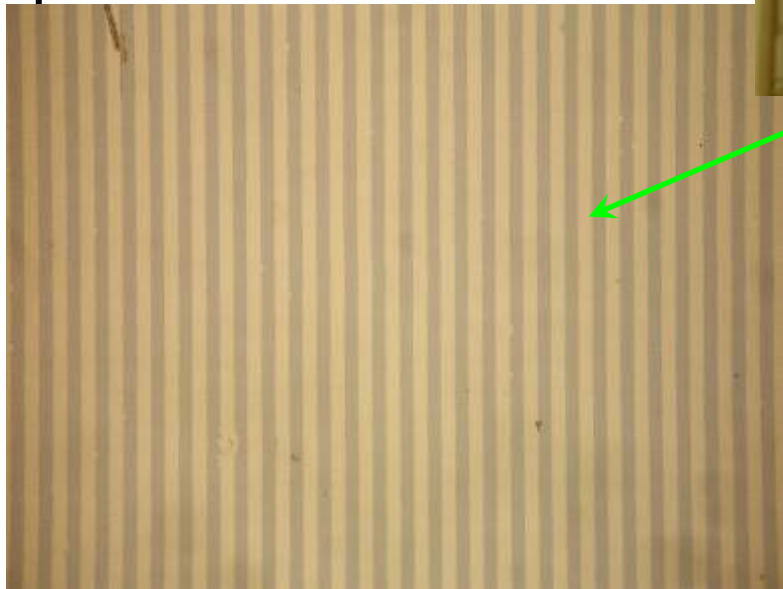
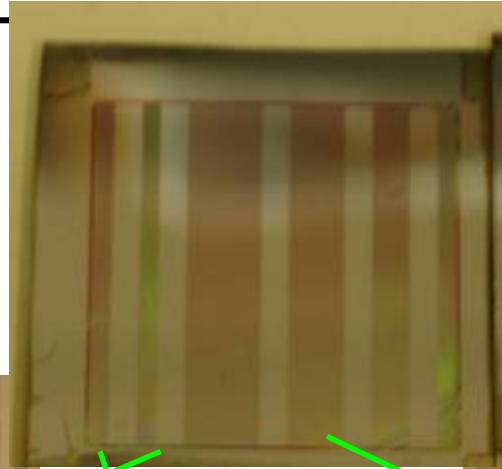
Vega ©Tescan
HARVEST



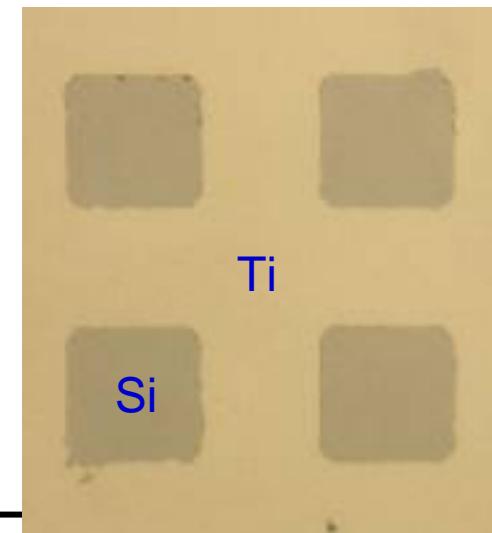
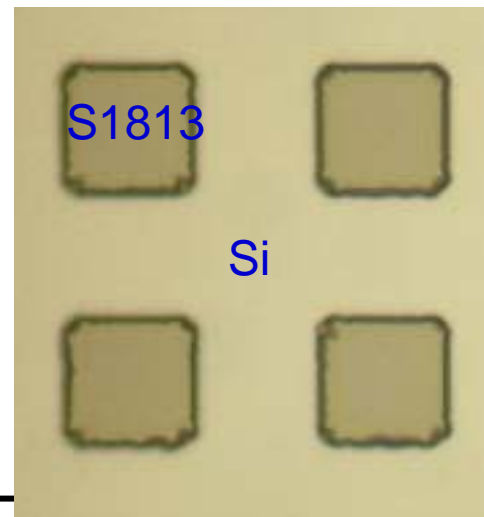
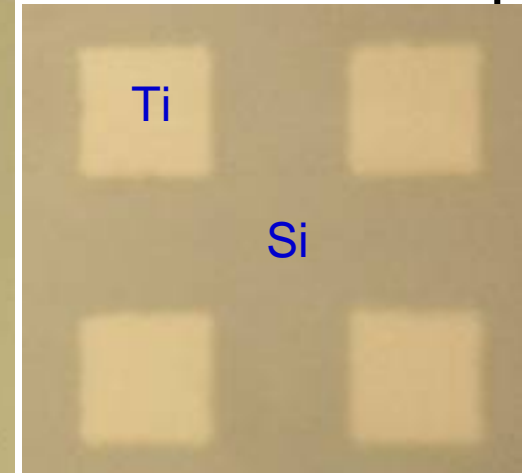
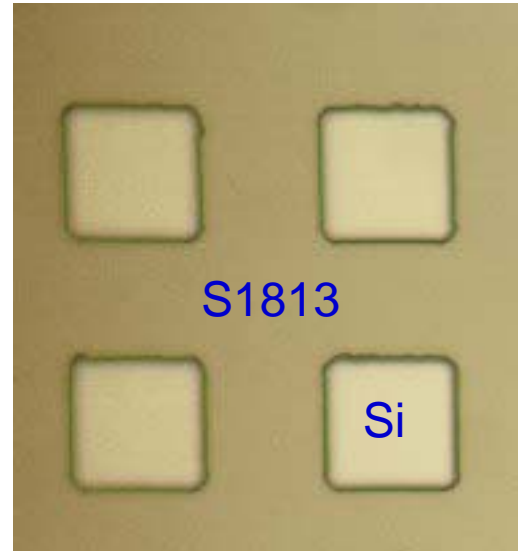
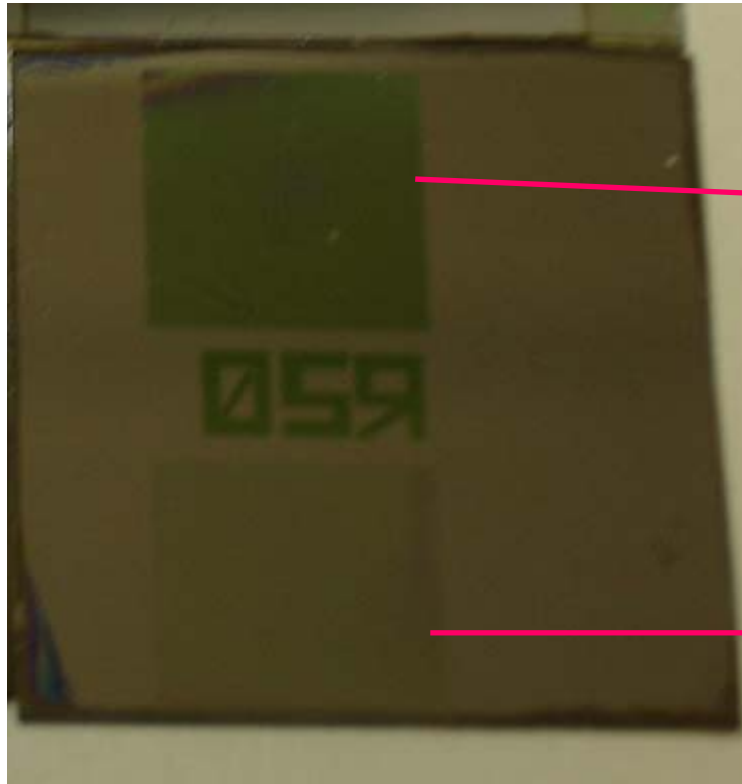
Lift OFF PROCESS BY ACETONE



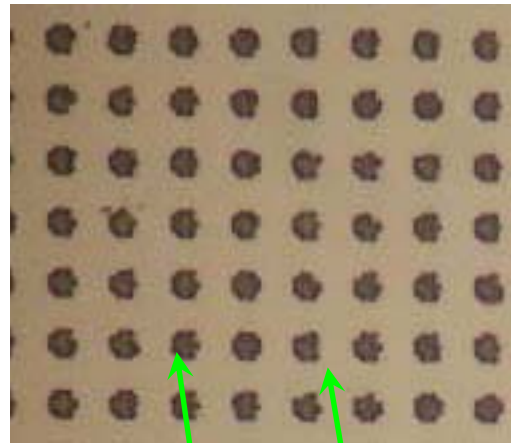
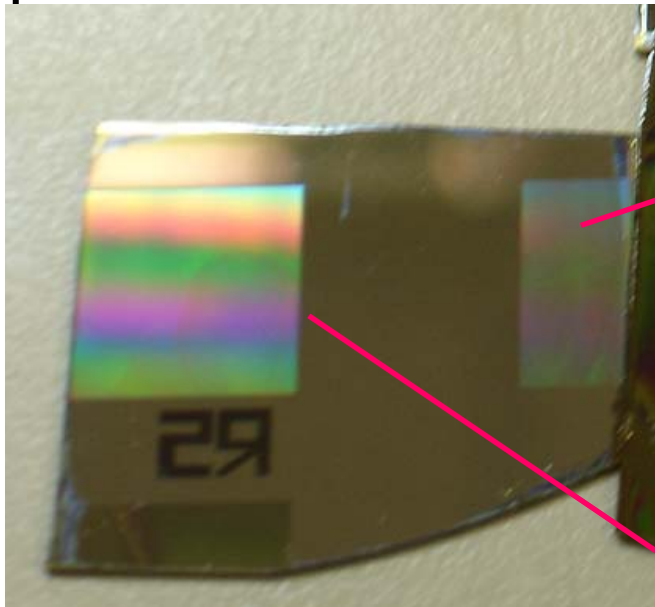
Ti 50 nm
LIFT OFF



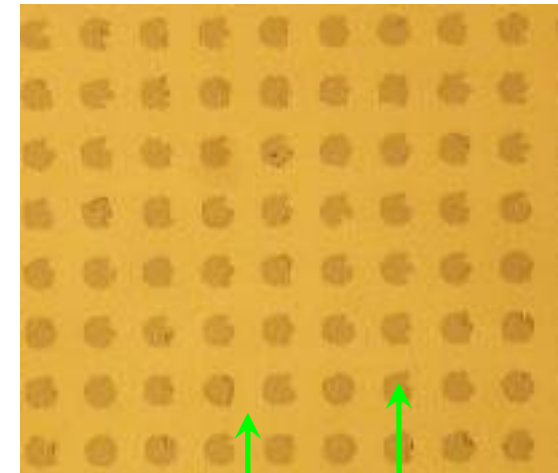
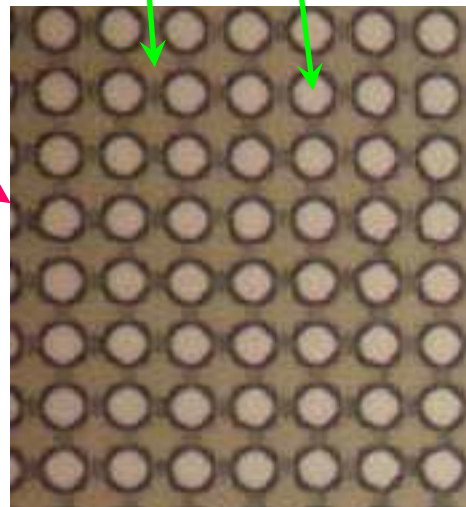
Ti 50 nm on Si 20 um



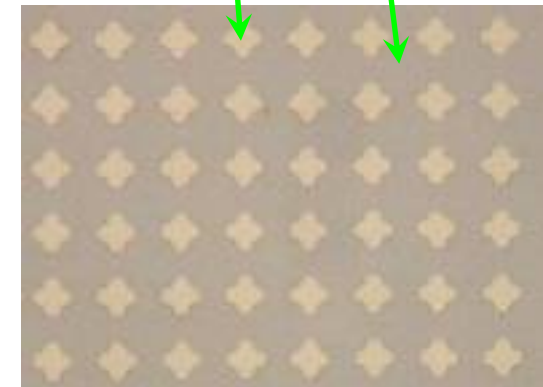
Ti 50 nm on Si 5 um



S1813 Si



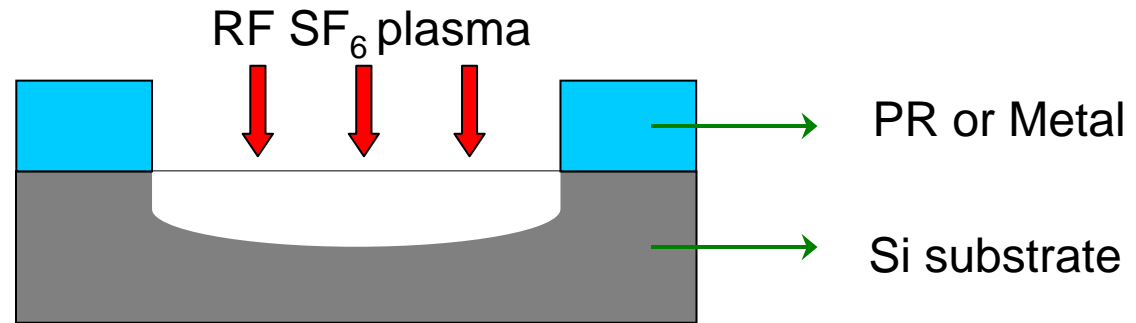
Ti Si



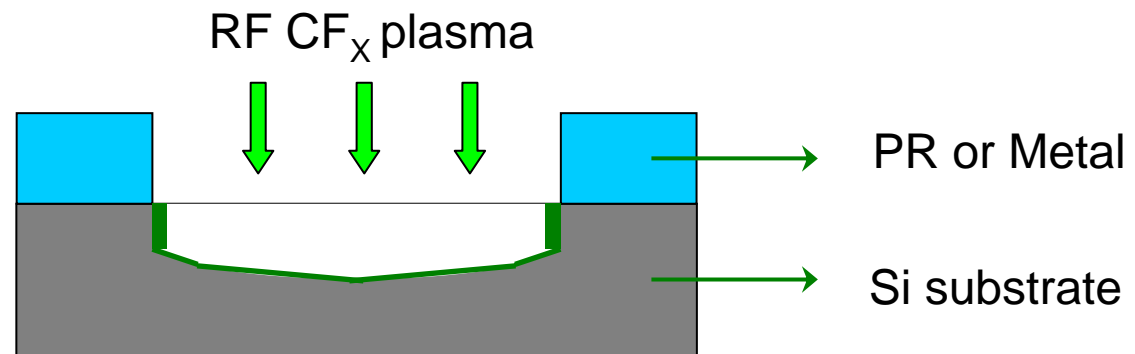
ICP- "BOSCH" Recipe

Etching & Sidewall Passivation Cycle

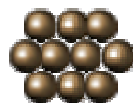
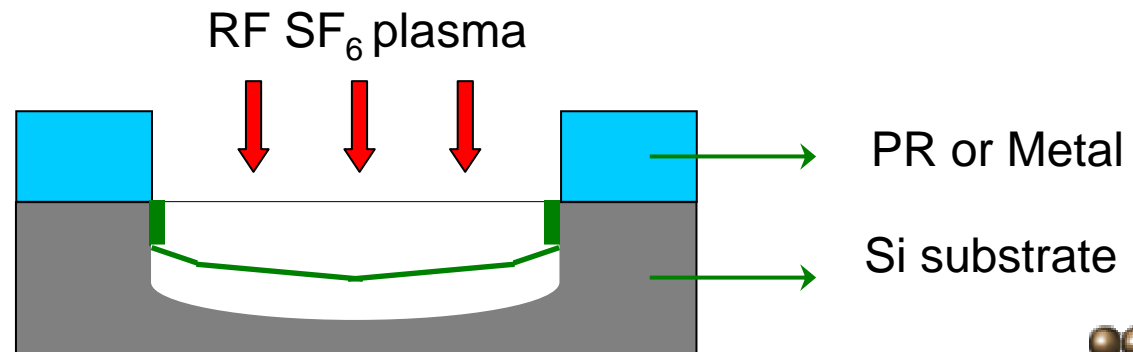
(a) Etch Step

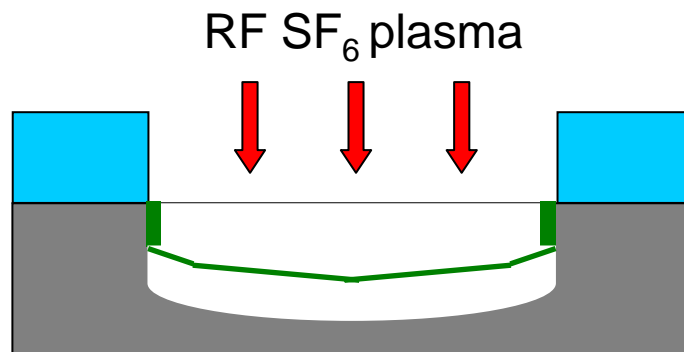
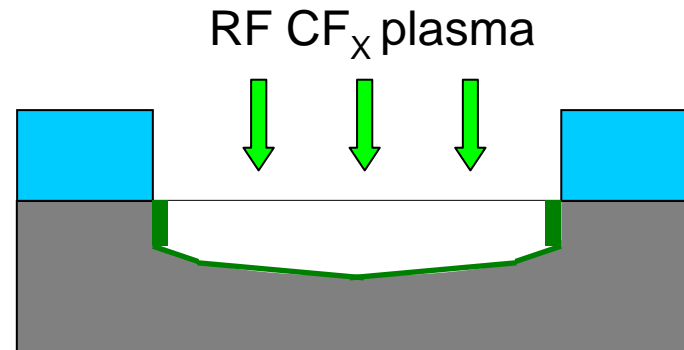
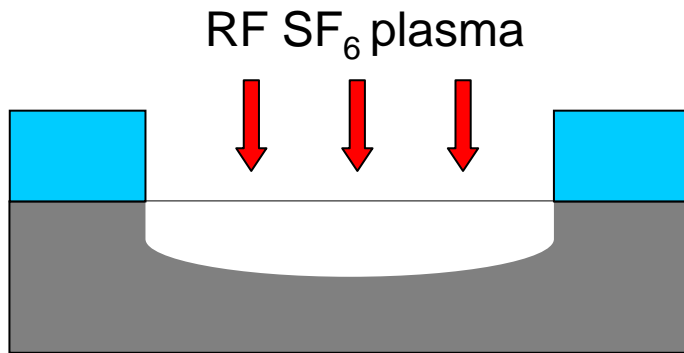


(b) Passivate Step



(a') Etch Step





The gases in RCAS

C_4F_8 , CF_4 , CHF_3 , Ar, O_2

SAMCO ICP (RECIPE)

Si etching $CF_4 / O_2 = 30 / 10$

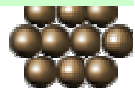
SiO_2 (on Si) $CHF_3 / Ar = 15 / 30$

The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

RECIPE

	TIME	
Etch:	11.5 s	$SF_6(130sccm)$ $O_2(13sccm)$
Passivate:	7s	$C_4F_8(85sccm)$



RCAS E-Beam Evaporator



一. 儀器名稱

中文名稱：電子束蒸鍍系統

英文名稱：E-Beam

二. 儀器廠牌、型號及儀器購置年限

廠牌：聚昌科技 AST

儀器購置年限：民國92年7月

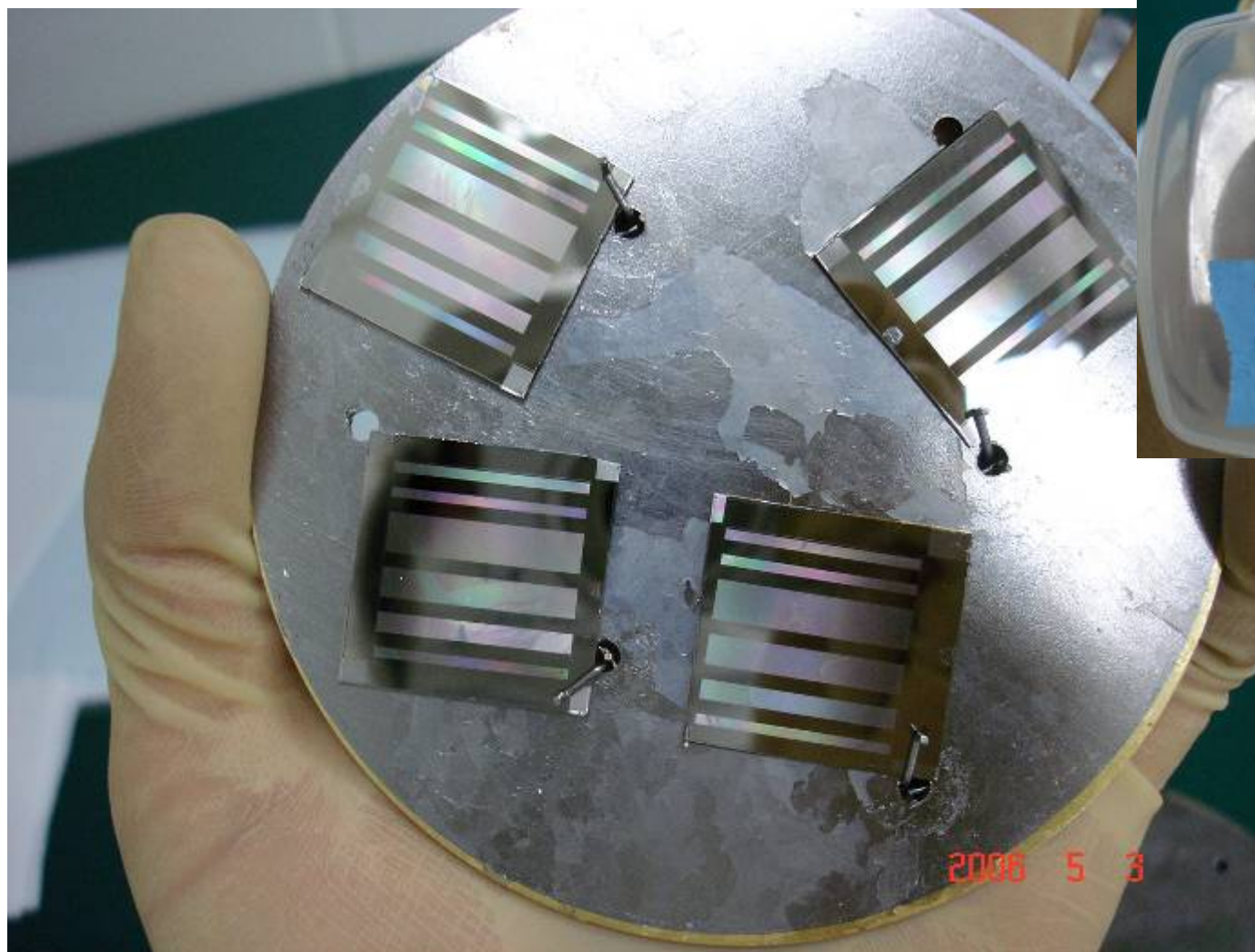
三. 重要規格

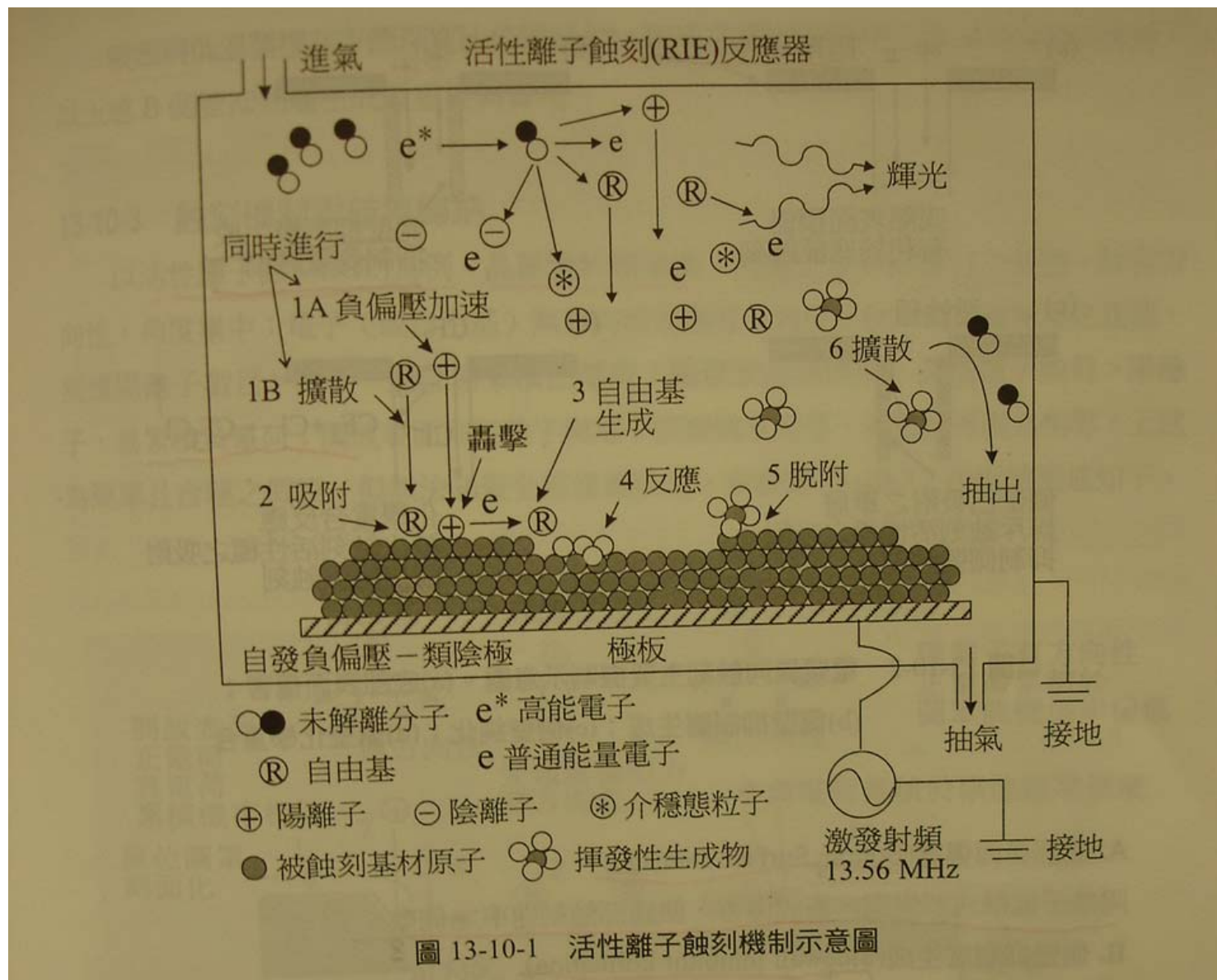
蒸鍍金屬: Ni、Ti、Au、Al、Pt、Cr

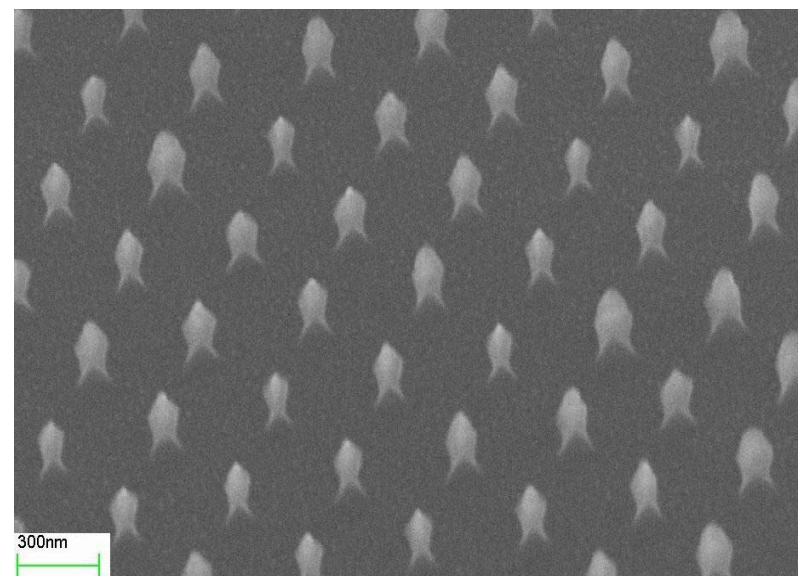
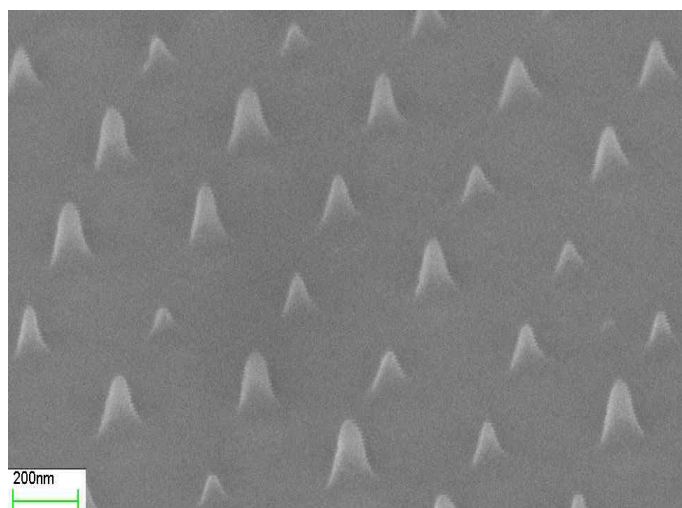
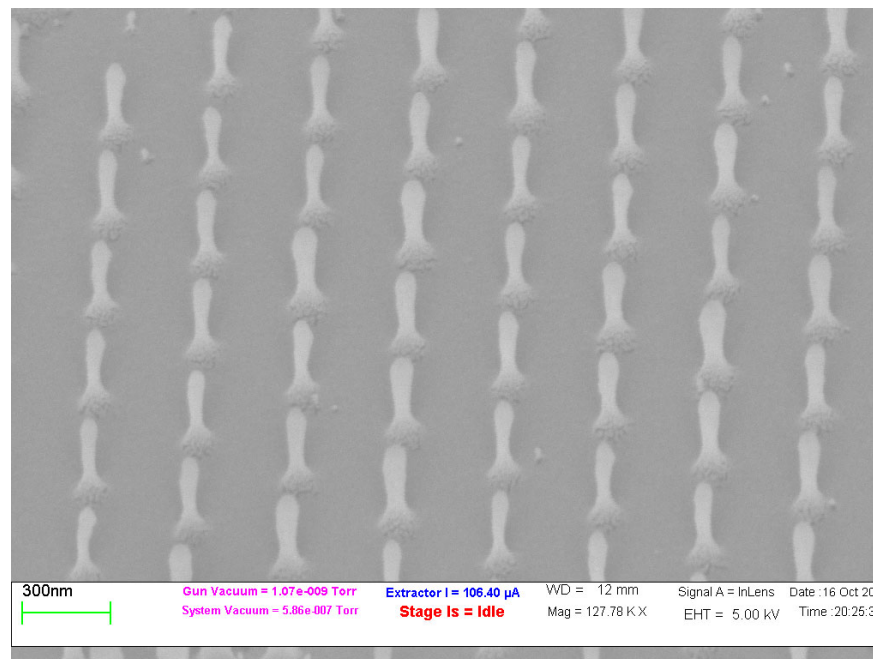
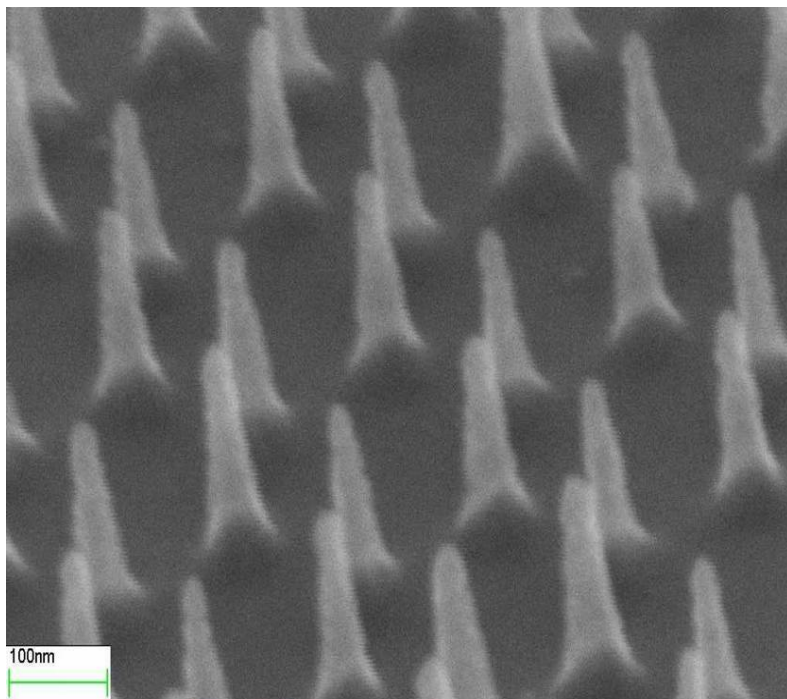
儀器配置圖



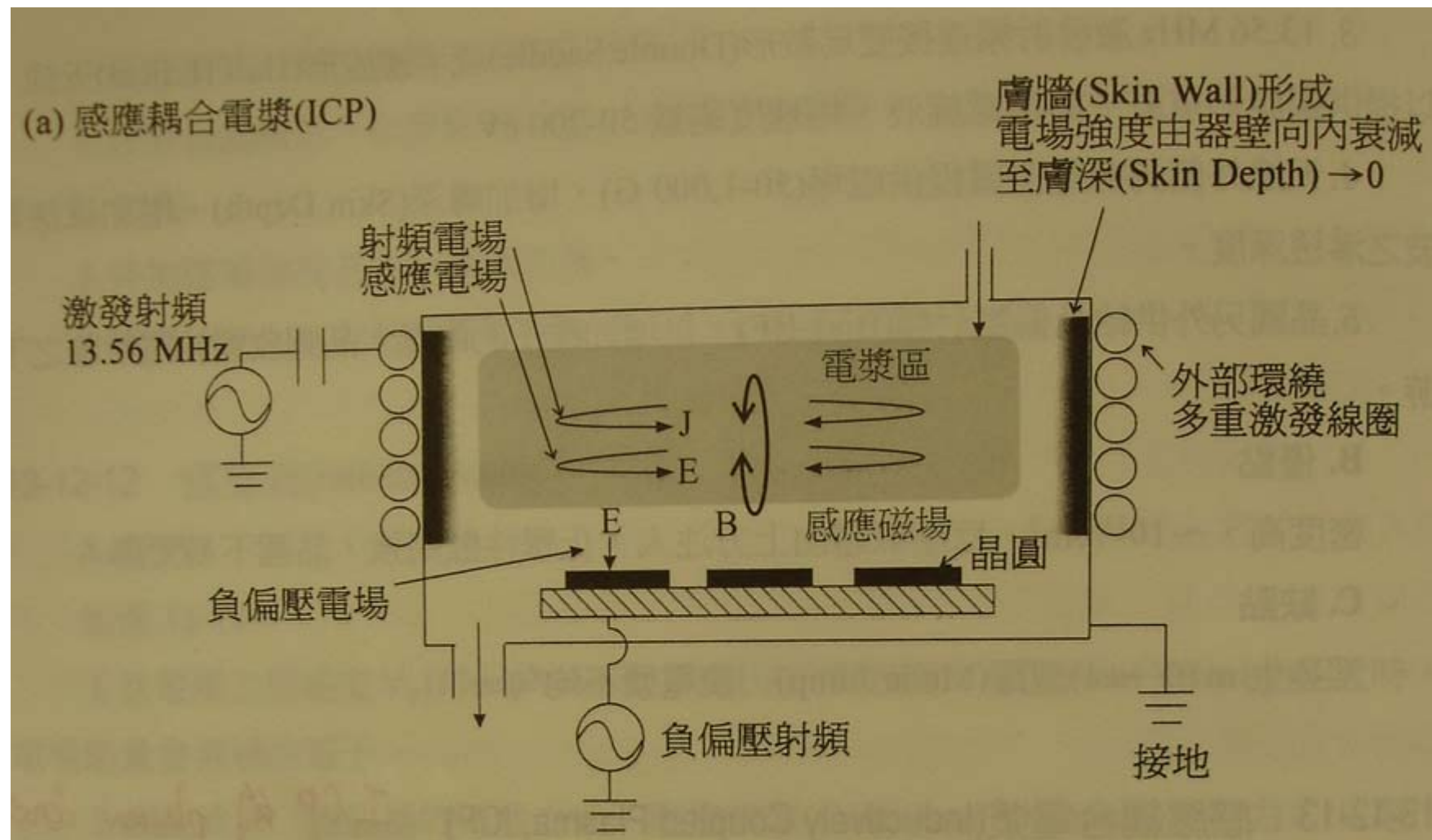
TI 50 nm



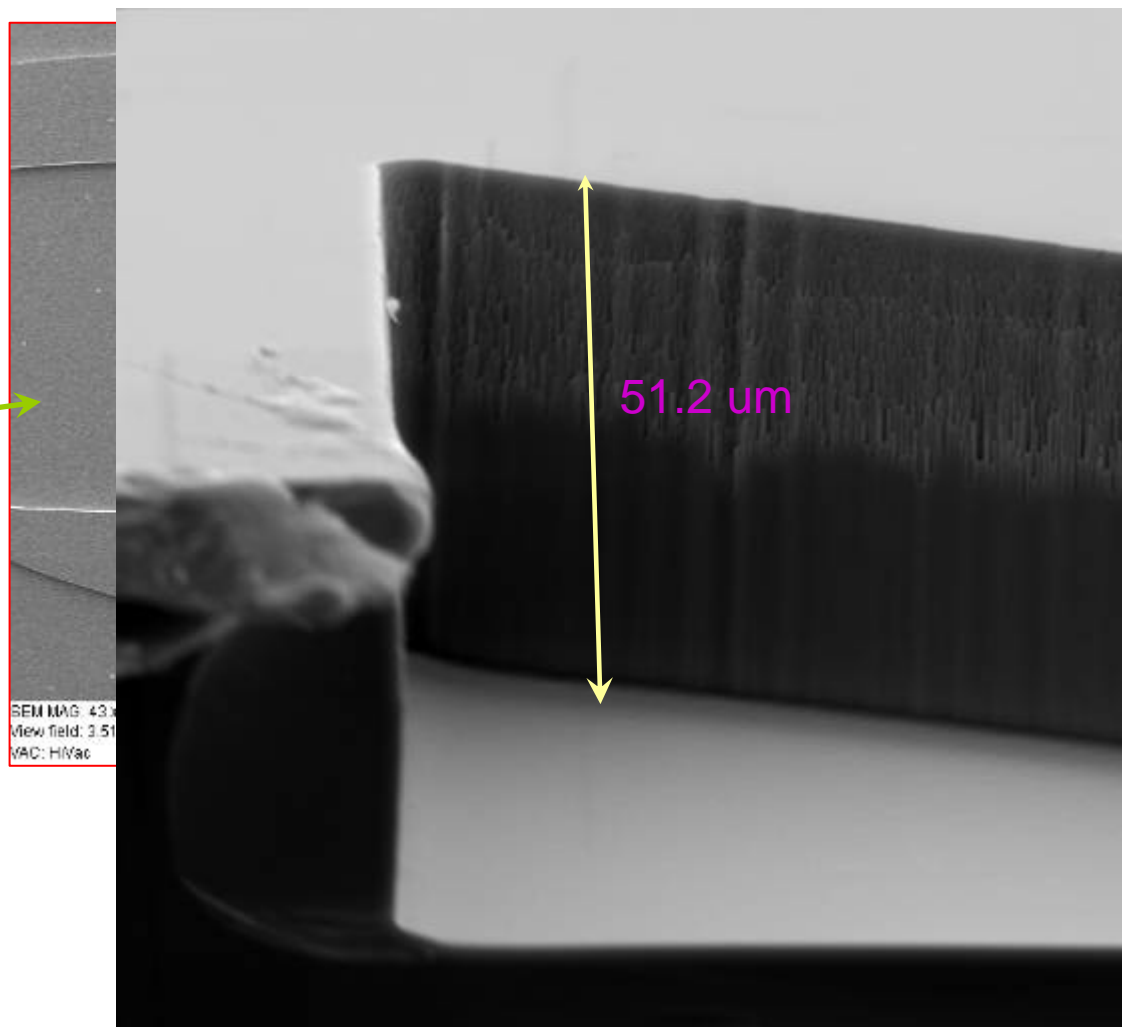




(a) 感應耦合電漿(ICP)



NEMSRC ICP



SEM MAG: 43.1 kx
View field: 3.51 um
VAC: HiVac

SEM MAG: 1.19 kx
View field: 126.40 um
VAC: HiVac

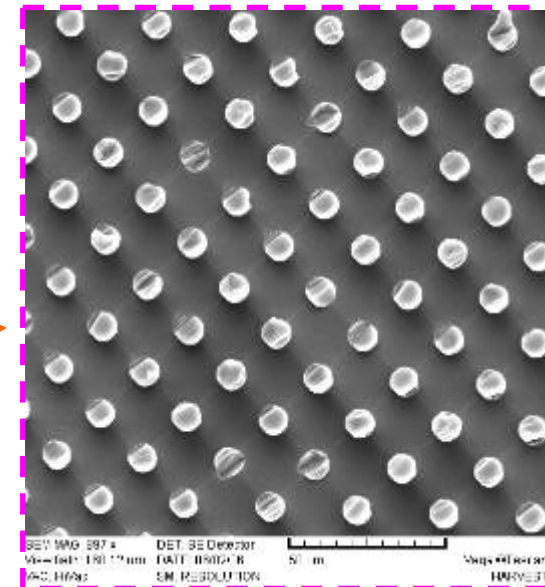
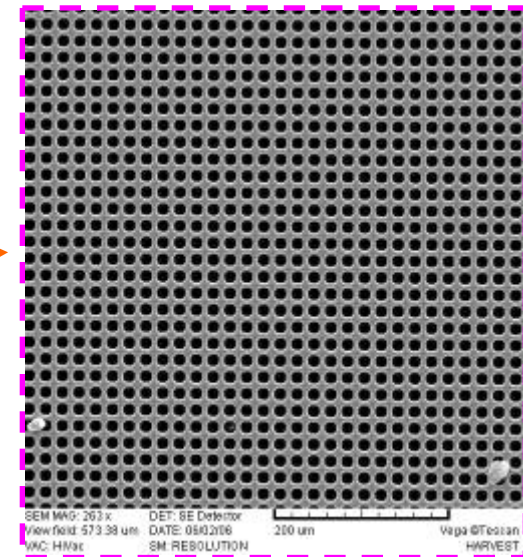
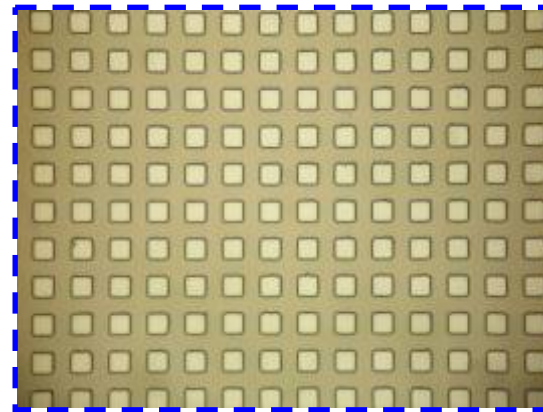
DET: SE Detector
DATE: 06/02/06
SM: RESOLUTION

50 um

Vega ©Tescan
HARVEST



Si ---S1813---ICP



NEMSRC ICP : S1813 on Si

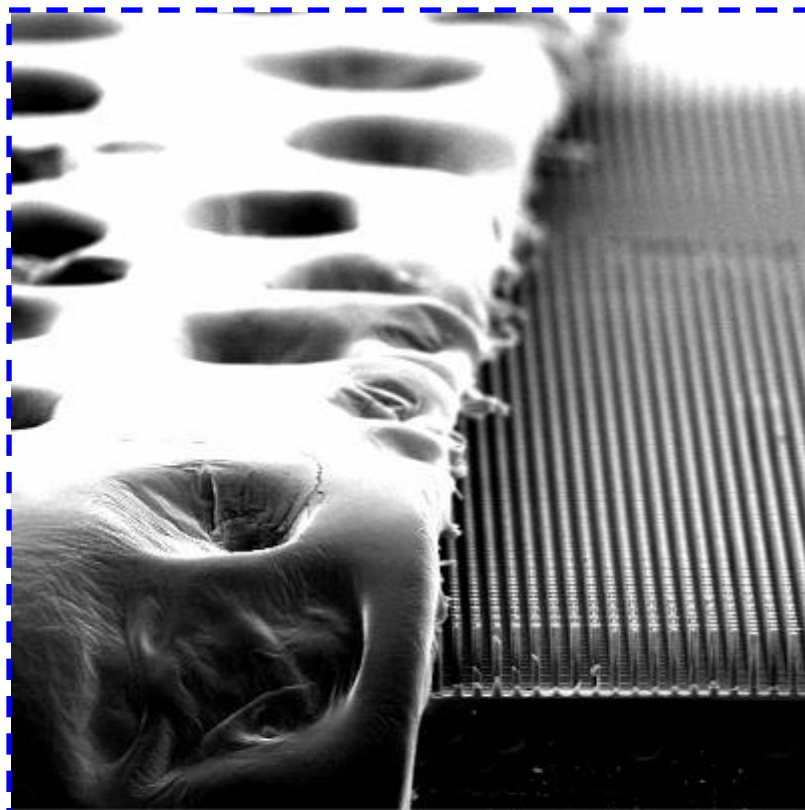
The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

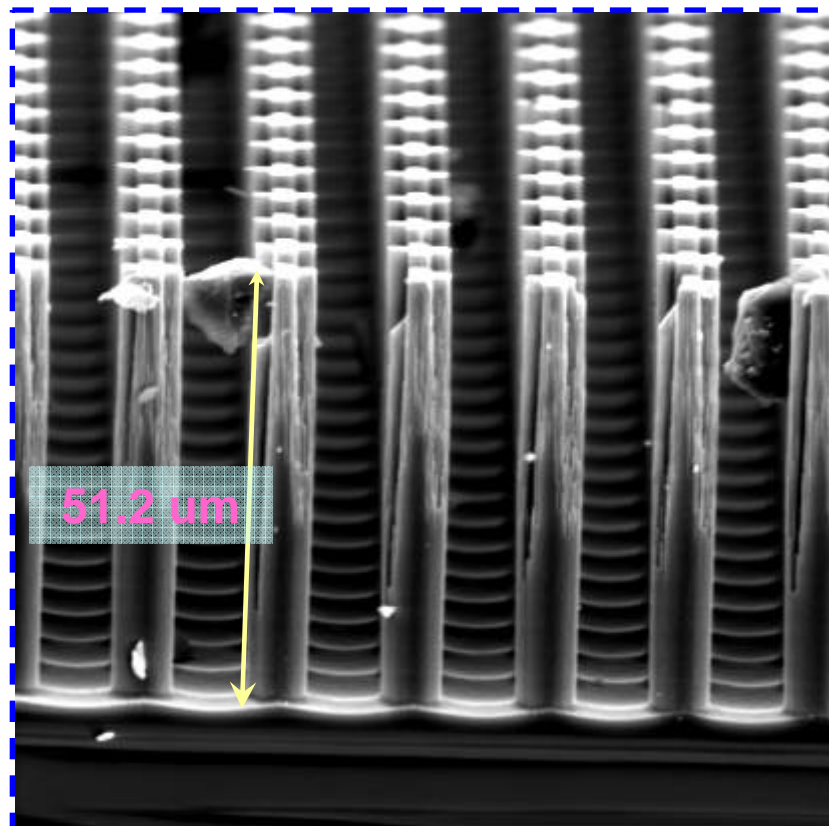
RECIPE

TIME

Etch: 11.5 s SF_6 (130sccm) O_2 (13sccm)
Passivate: 7s C_4F_8 (85sccm)



SEM MAG: 191 x DET: SE Detector
View field: 789.53 um DATE: 06/02/06
VAC: HiVac SM: RESOLUTION
Vega ©Tescan
HARVEST



SEM MAG: 1,28 kx DET: SE Detector
View field: 117.35 um DATE: 06/02/06
VAC: HiVac SM: RESOLUTION
Vega ©Tescan
HARVEST

NEMSRC ICP : S1813 on Si --20 μ m array

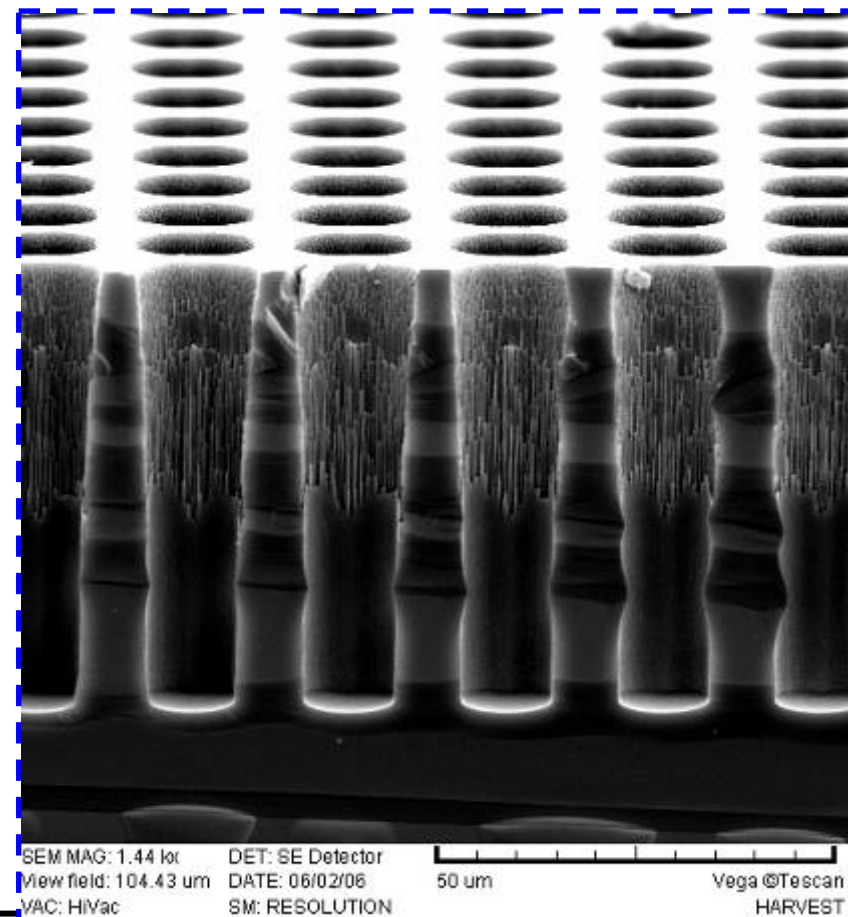
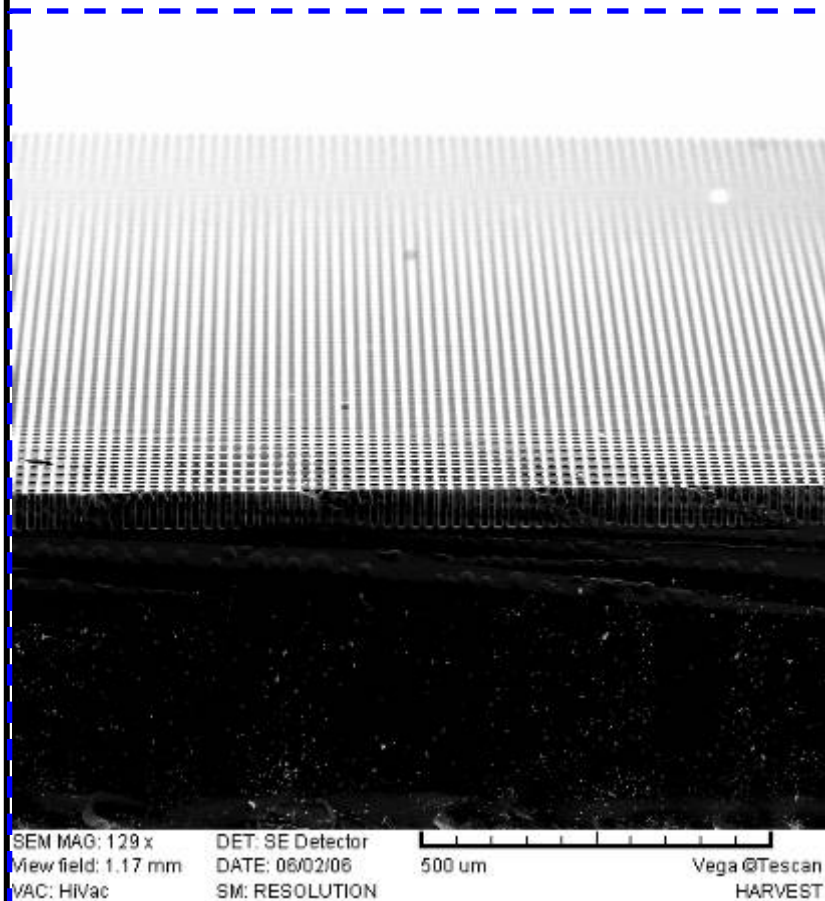
The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

RECIPE

TIME

Etch: 11.5 s SF_6 (130sccm) O_2 (13sccm)
Passivate: 7s C_4F_8 (85sccm)



NEMSRC ICP : S1813 on Si --20 μ m array

The gases in NEMSRC

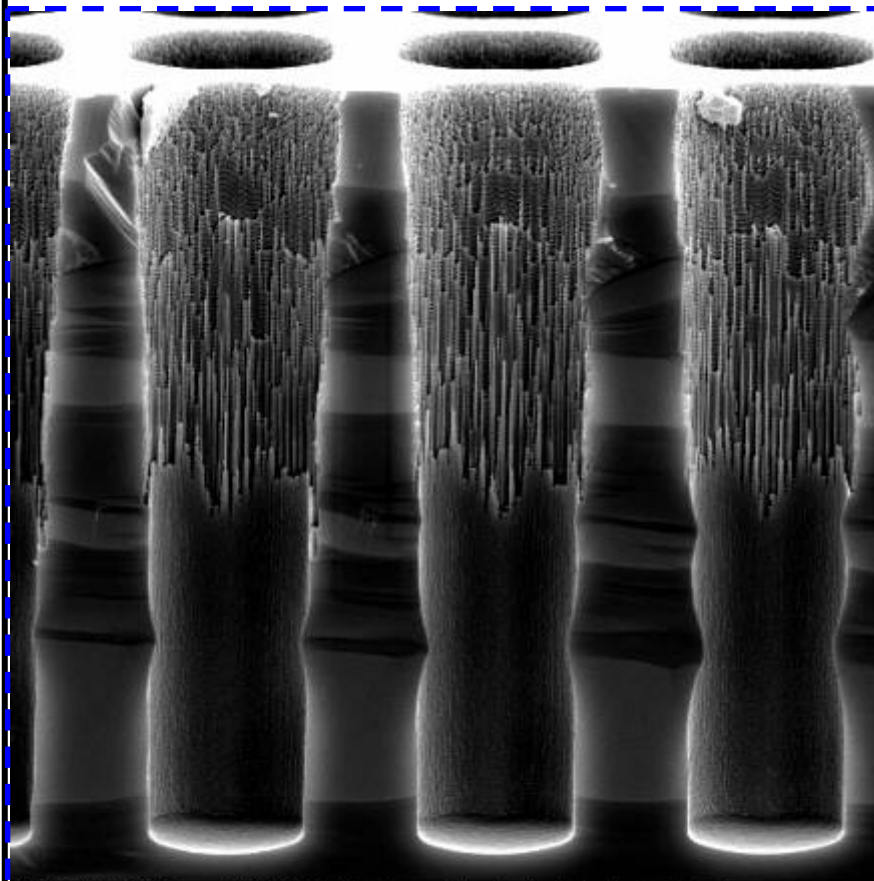
SF_6 , C_4F_8 , CF_4 , O_2

RECIPE

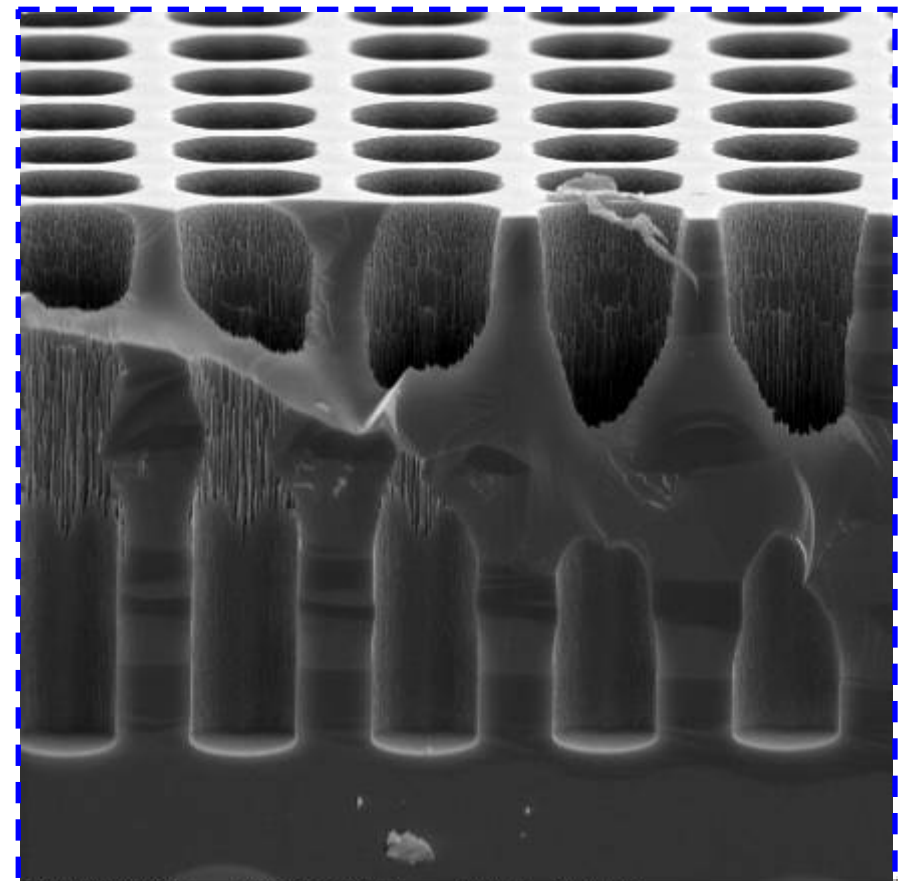
TIME

Etch: 11.5 s SF_6 (130sccm) O_2 (13sccm)

Passivate: 7s C_4F_8 (85sccm)



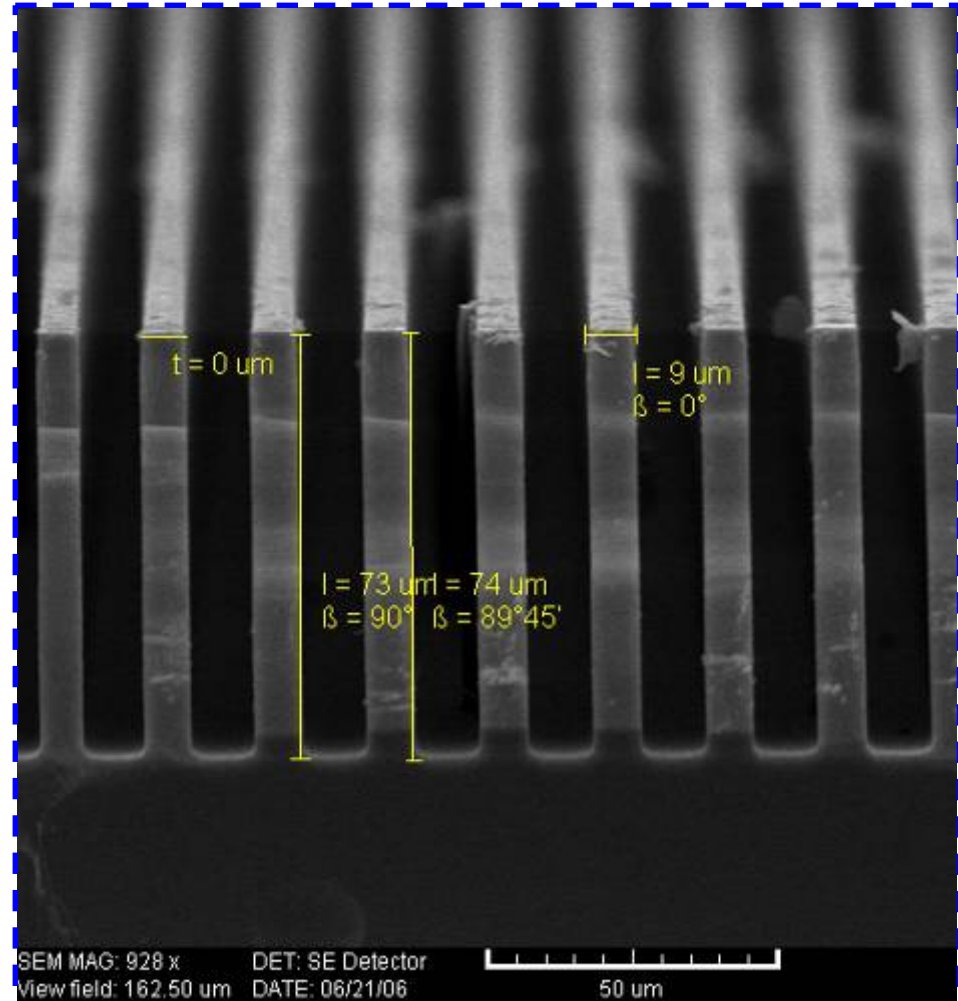
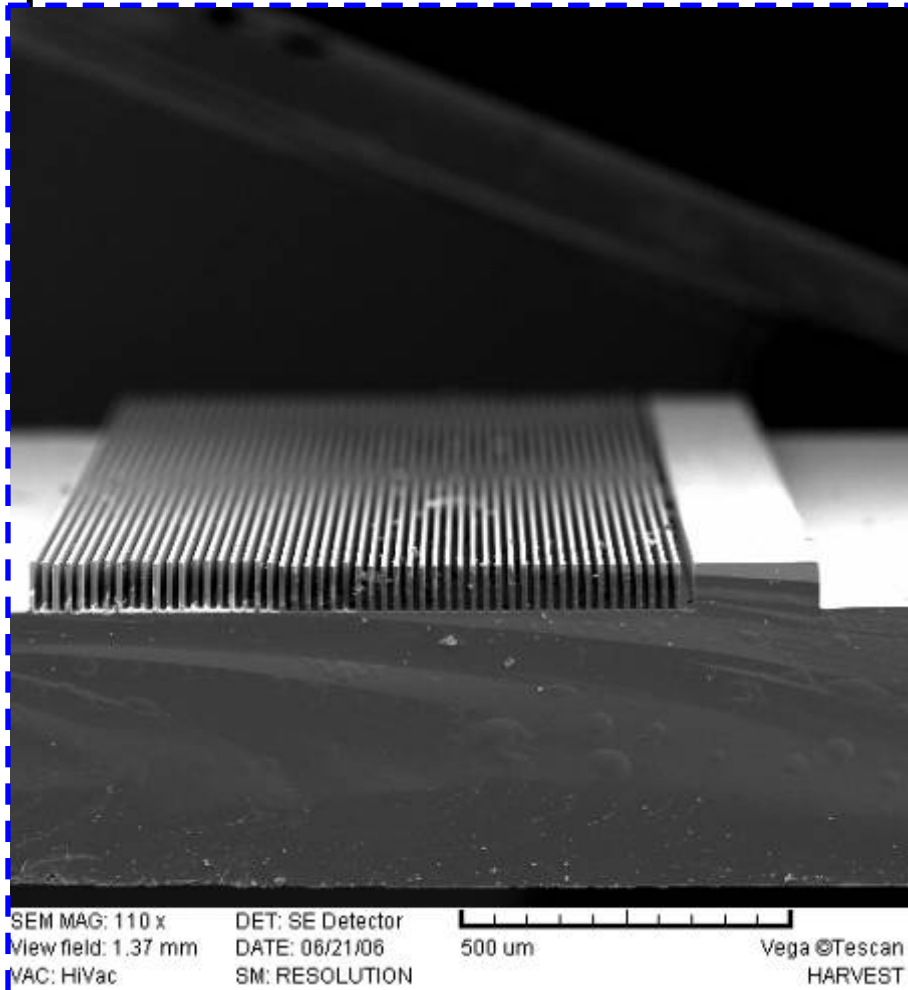
SEM MAG: 2.39 kx DET: SE Detector
View field: 63.18 μ m DATE: 06/02/06
VAC: HiVac SM: RESOLUTION
20 μ m
Vega ©Tescan
HARVEST



SEM MAG: 1.60 kx DET: SE Detector
View field: 94.37 μ m DATE: 06/02/06
VAC: HiVac SM: RESOLUTION
20 μ m
Vega ©Tescan
HARVEST

NEMSRC ICP

Si 10 μm Line



NEMSRC ICP : S1813 on Si

10 um pillar

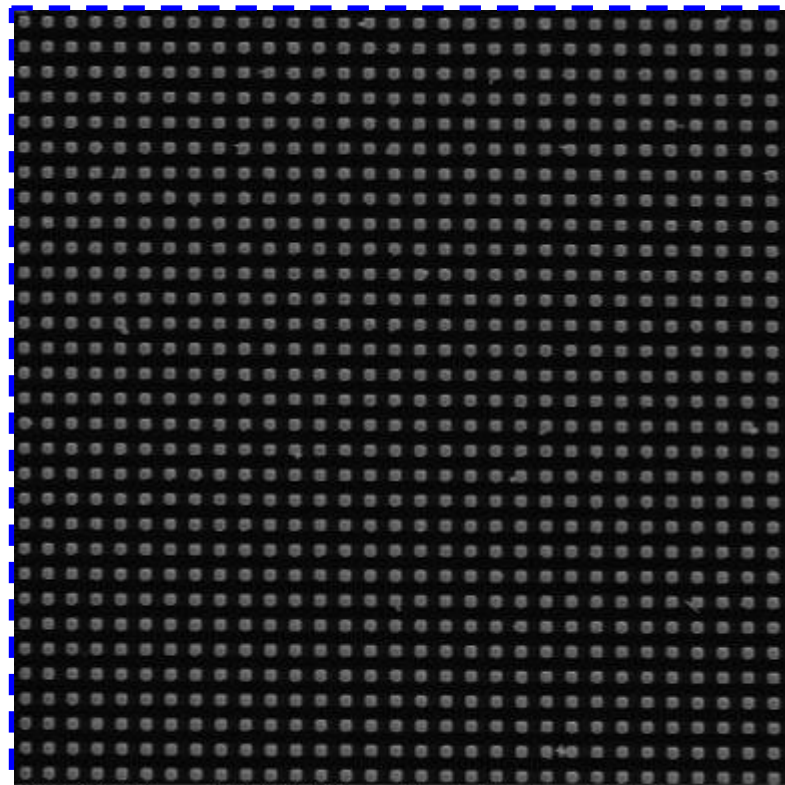
The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

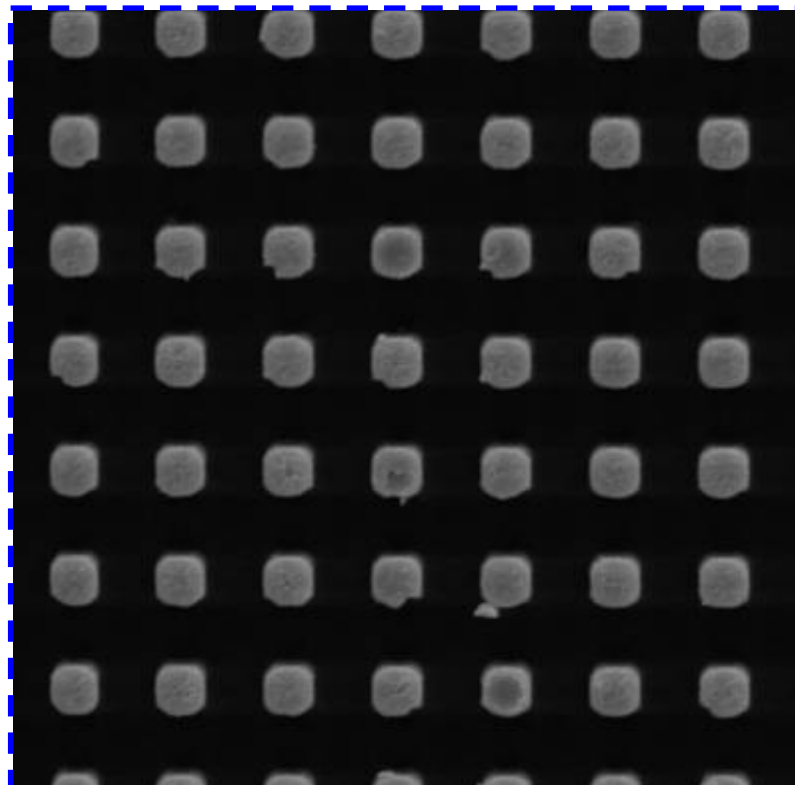
RECIPE

TIME

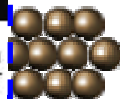
Etch: 11.5 s SF_6 (130sccm) O_2 (13sccm)
Passivate: 7s C_4F_8 (85sccm)



SEM MAG: 251 x DET: SE Detector
View field: 600.60 um DATE: 06/21/08 200 um Vega @Tescan
VAC: HiVac SM: RESOLUTION HARVEST

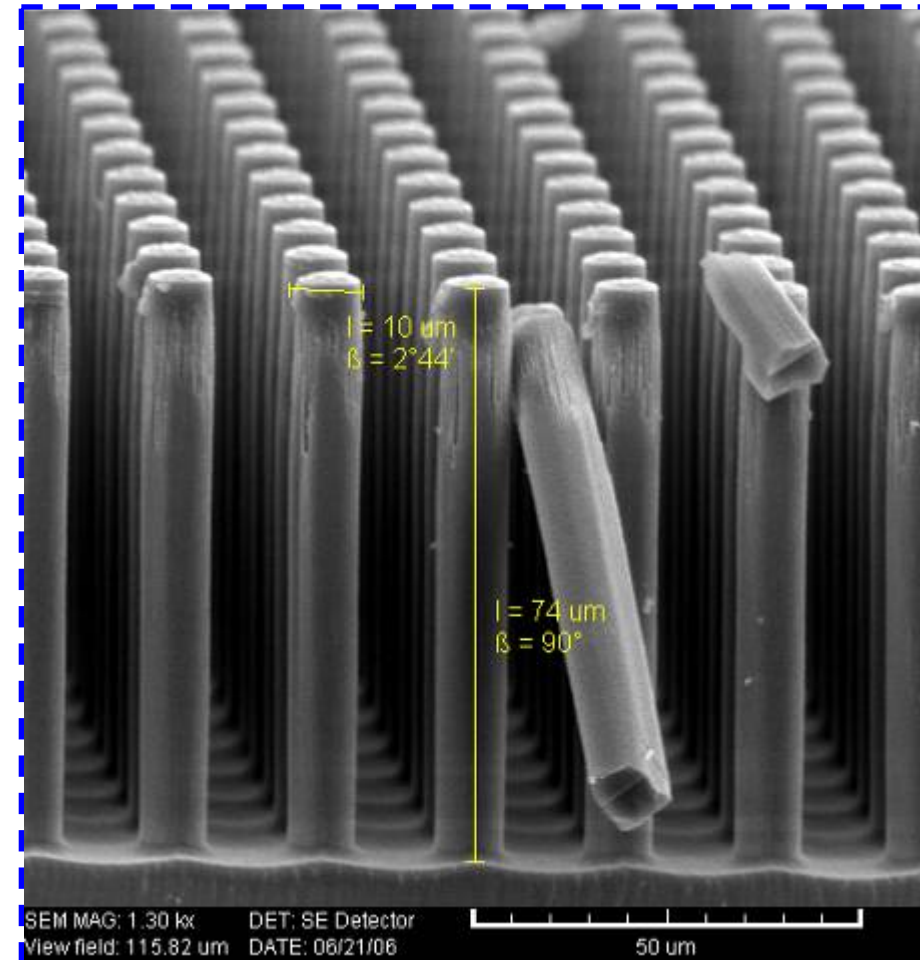
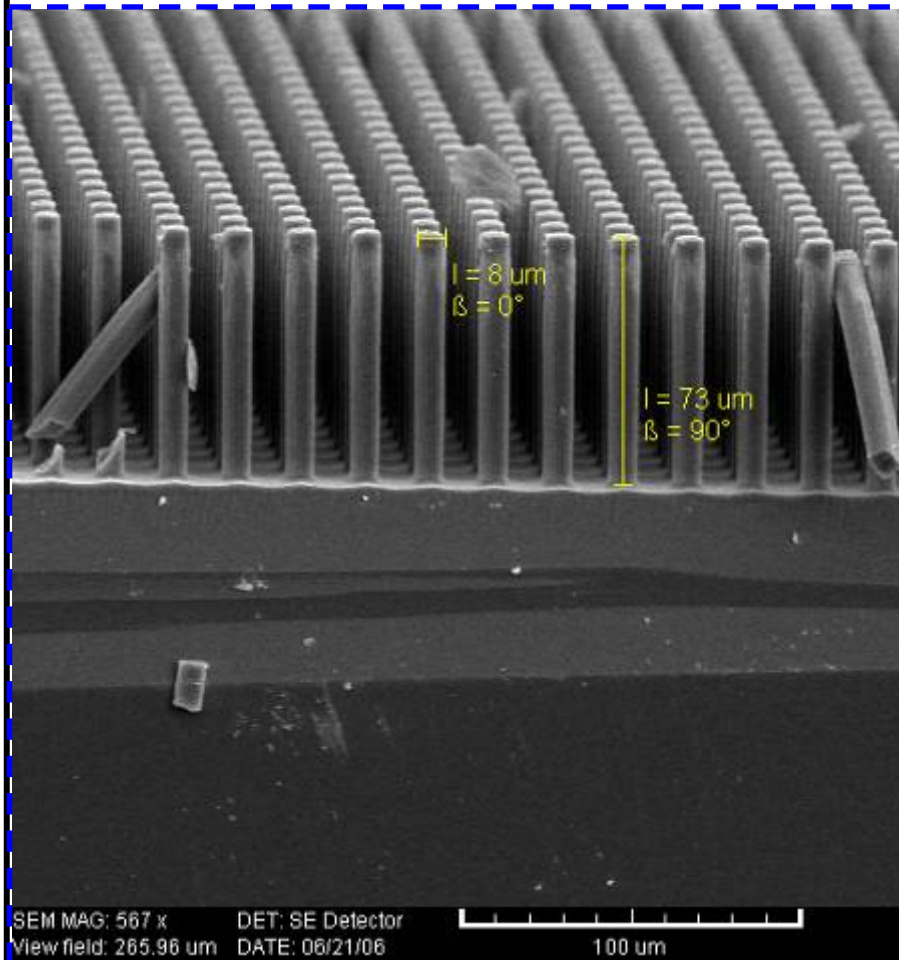


SEM MAG: 1.07 kc DET: SE Detector
View field: 141.20 um DATE: 06/21/08 50 um Vega @Tescan
VAC: HiVac SM: RESOLUTION HARVEST

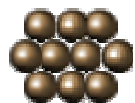
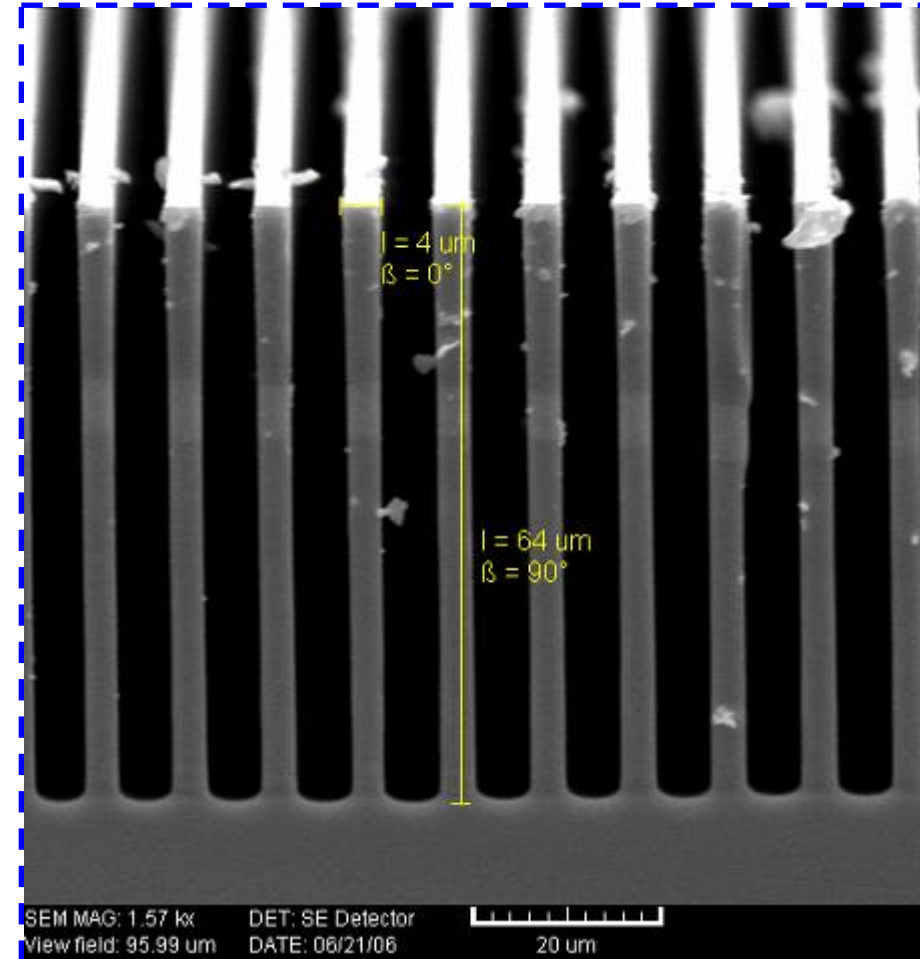
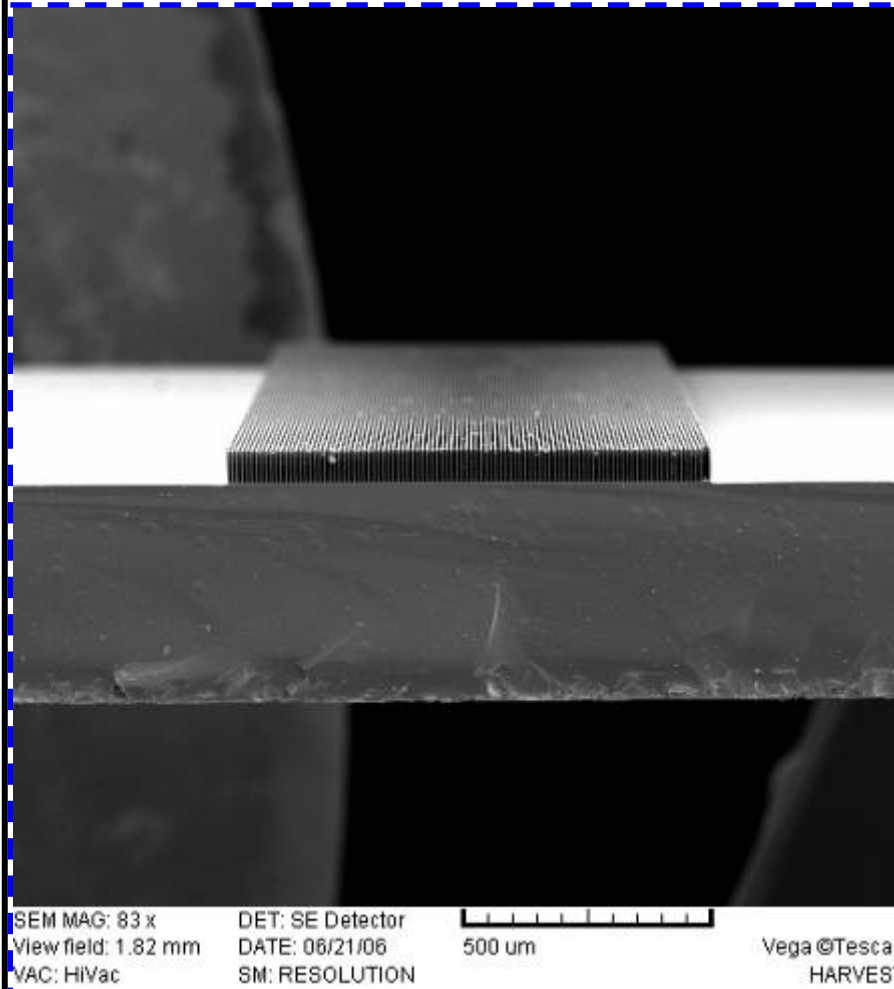


NEMSRC ICP

Si 10 μm pillar



Si 5um Line



NEMSRC ICP : S1813 on Si

5 um pillar

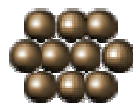
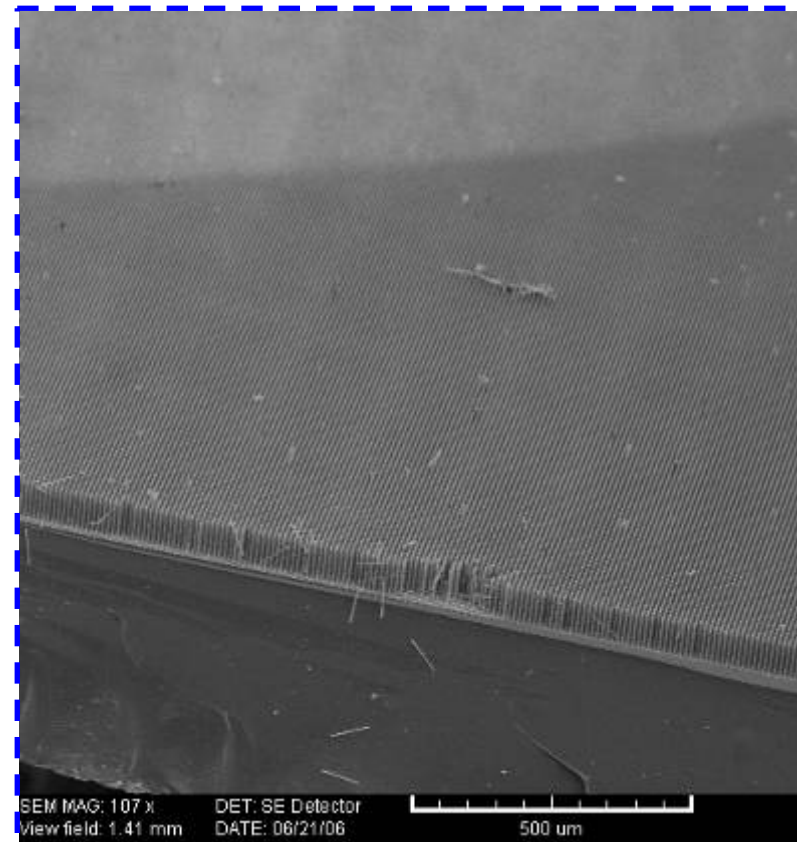
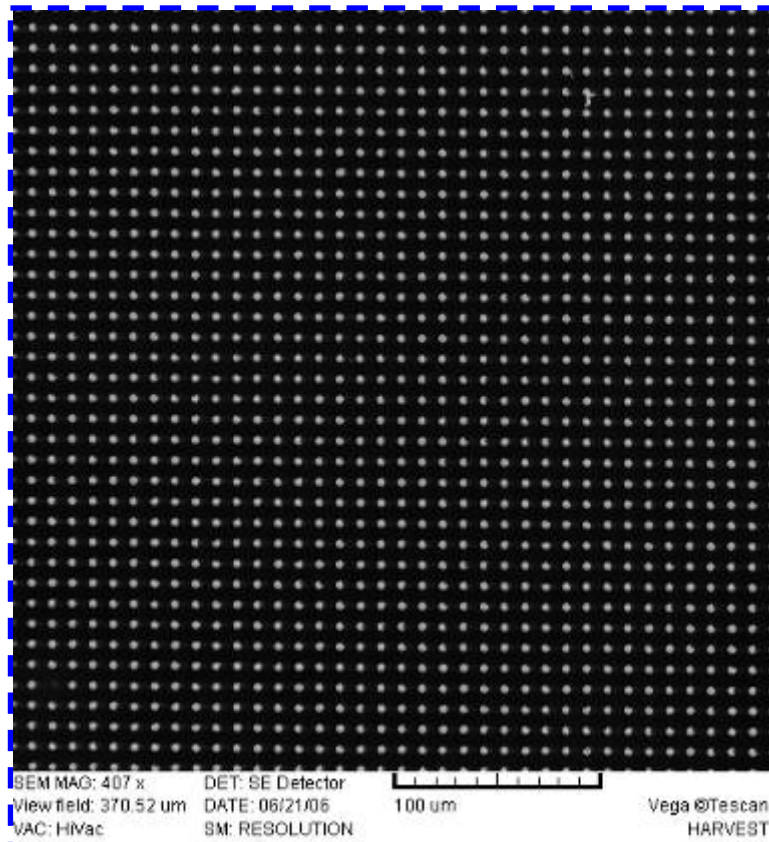
The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

RECIPE

TIME

Etch: 11.5 s SF_6 (130sccm) O_2 (13sccm)
Passivate: 7s C_4F_8 (85sccm)



NEMSRC ICP : S1813 on Si

5 um pillar

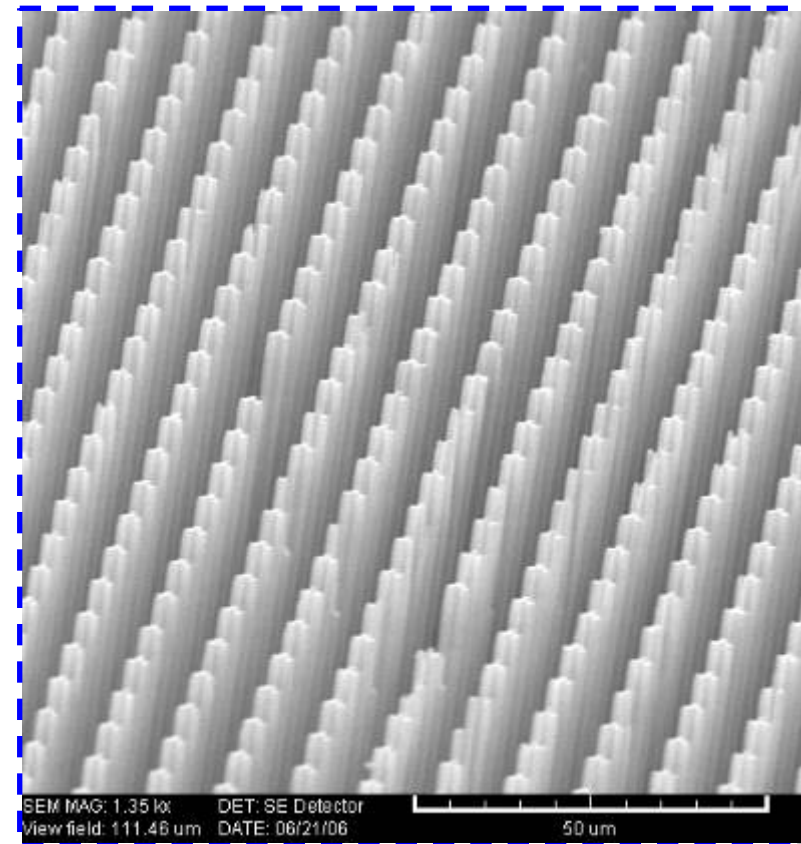
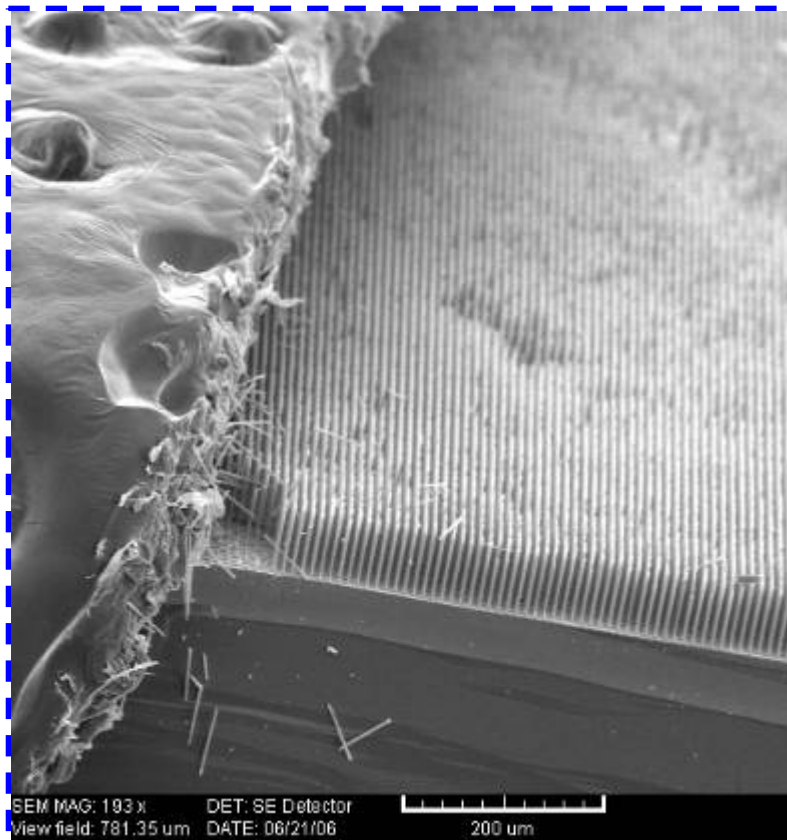
The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

RECIPE

TIME

Etch: 11.5 s	SF_6 (130sccm)	O_2 (13sccm)
Passivate: 7s	C_4F_8 (85sccm)	



NEMSRC ICP : S1813 on Si

5 um pillar

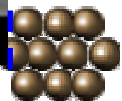
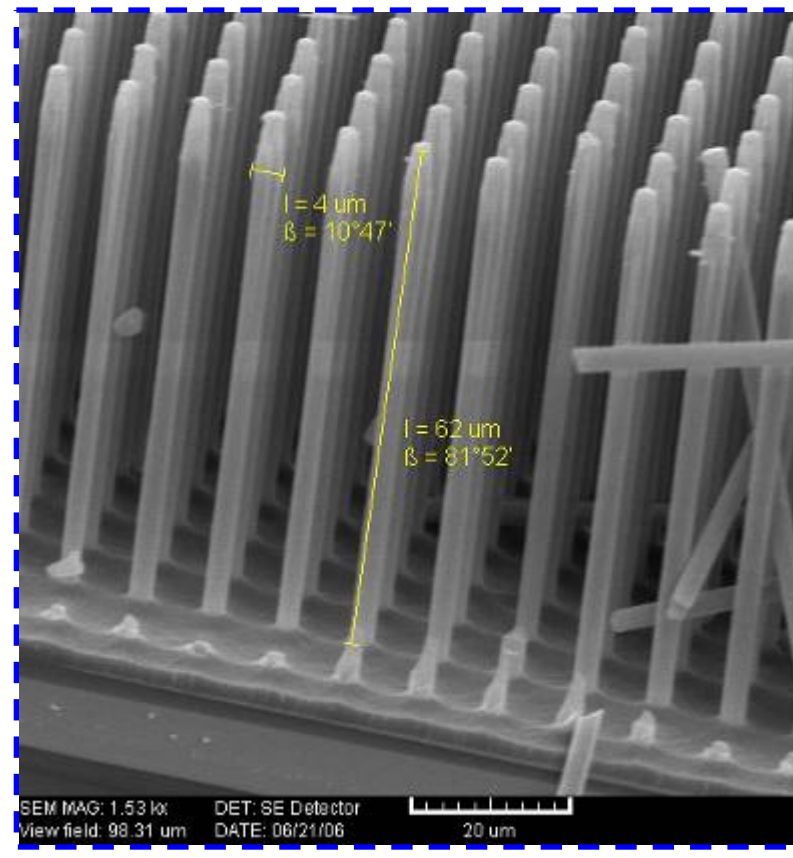
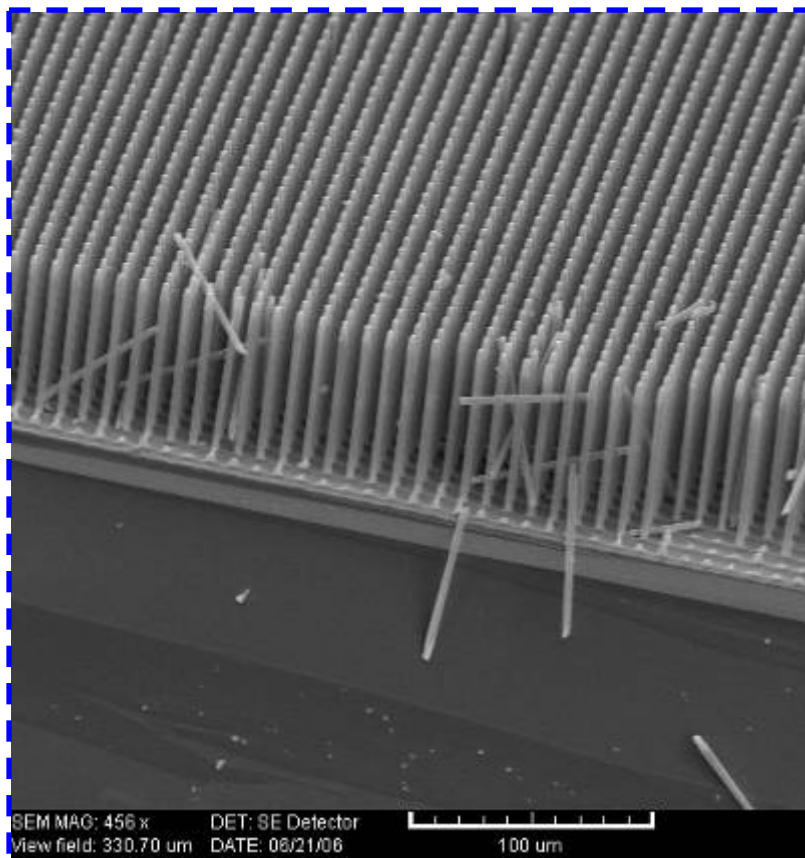
The gases in NEMSRC

SF_6 , C_4F_8 , CF_4 , O_2

RECIPE

TIME

Etch: 11.5 s $\text{SF}_6(130\text{sccm})$ $\text{O}_2(13\text{sccm})$
Passivate: 7s $\text{C}_4\text{F}_8(85\text{sccm})$



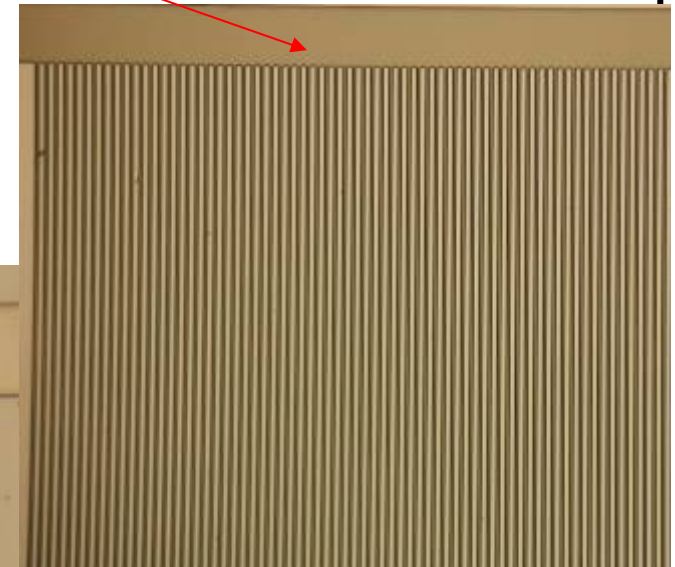
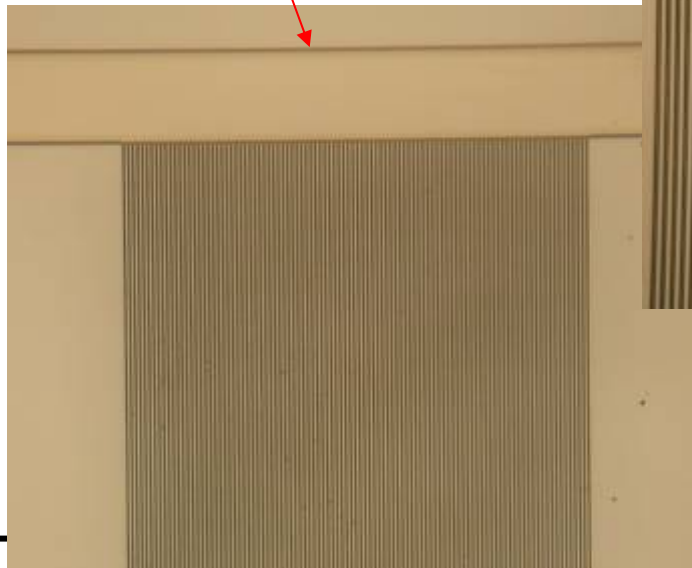
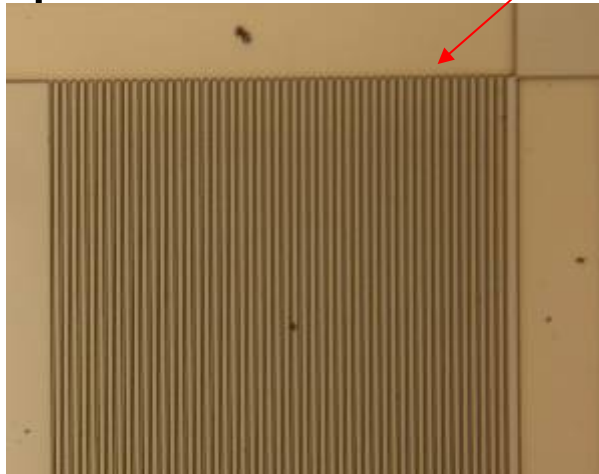
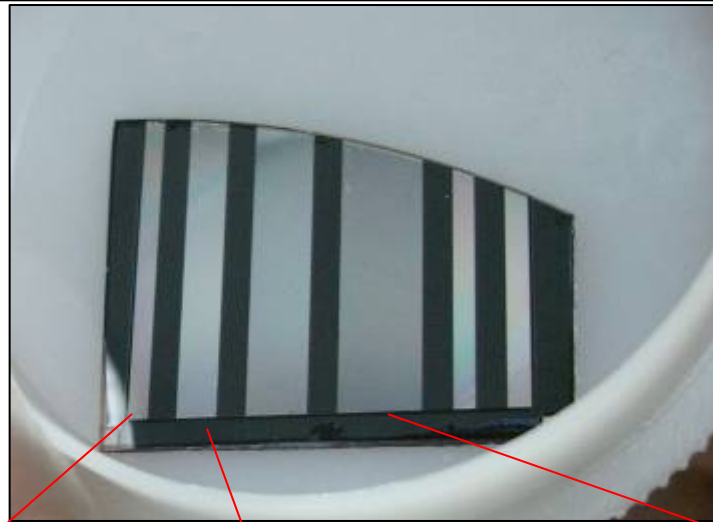
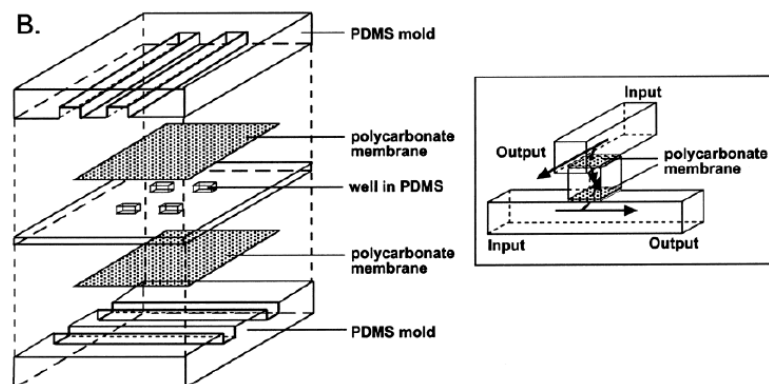
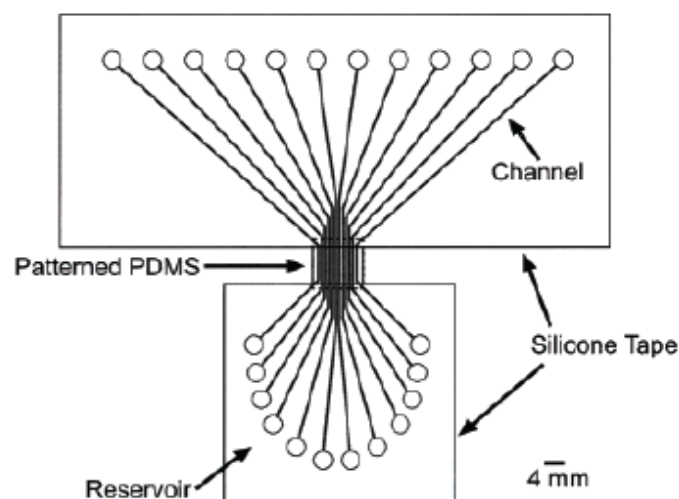


Table 1. Physical and Chemical Properties of PDMS

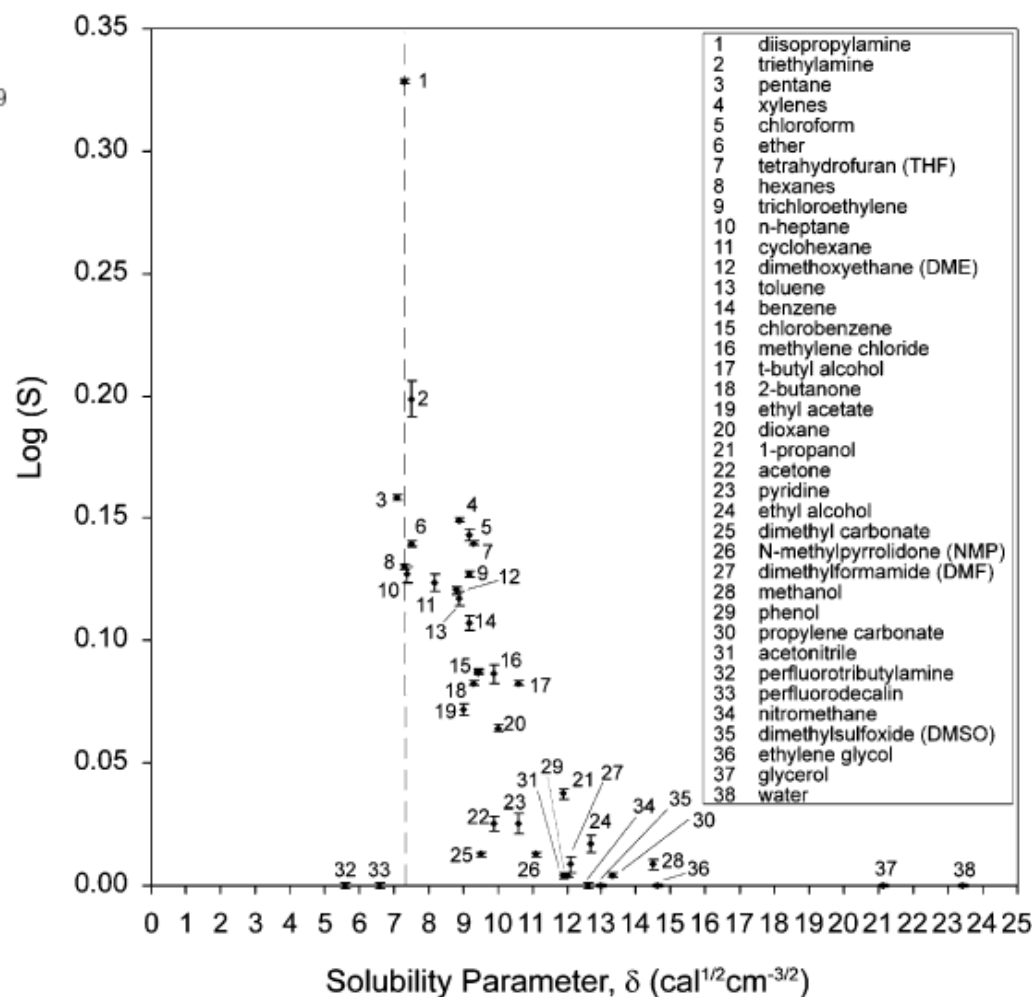
property	characteristic	consequence
optical	transparent; UV cutoff, 240 nm	optical detection from 240 to 1100 nm
electrical	insulating; breakdown voltage, 2×10^7 V/m ⁷¹	allows embedded circuits; intentional breakdown to open connections ⁴³
mechanical	elastomeric; tunable Young's modulus, typical value of ~ 750 kPa ²⁴	conforms to surfaces; allows actuation by reversible deformation; ²⁴ facilitates release from molds
thermal	insulating; thermal conductivity, 0.2 W/(m·K); coefficient of thermal expansion, $310 \mu\text{m}/(\text{m}\cdot^\circ\text{C})$ ⁷¹	can be used to insulate heated solutions; ⁶⁴ does not allow dissipation of resistive heating from electrophoretic separation
interfacial	low surface free energy ~ 20 erg/cm ² ²⁰	replicas release easily from molds; can be reversibly sealed to materials
permeability	impermeable to liquid water; permeable to gases and nonpolar organic solvents	contains aqueous solutions in channels; allows gas transport through the bulk material; incompatible with many organic solvents
reactivity	inert; can be oxidized by exposure to a plasma; $\text{Bu}_4\text{N}^+\text{F}^-$ (TBA)F	unreactive toward most reagents; surface can be etched; can be modified to be hydrophilic and also reactive toward silanes; ²⁰ etching with (TBA)F can alter topography of surfaces ⁵⁹
toxicity	nontoxic.	can be implanted in vivo; supports mammalian cell growth ^{57,59}



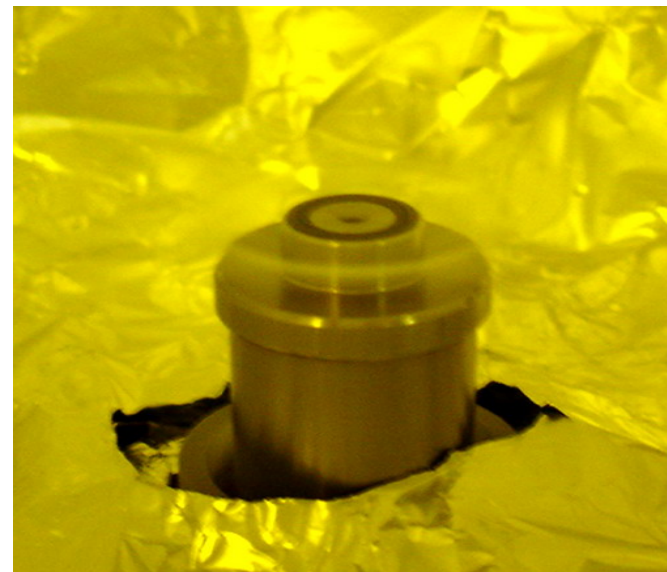
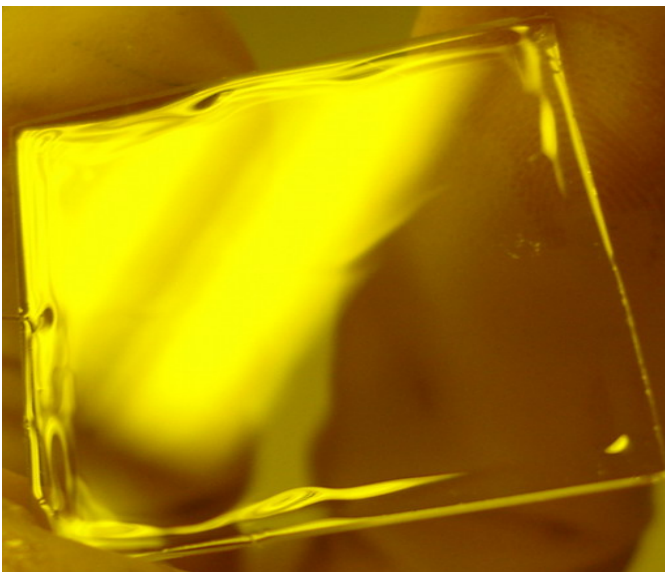
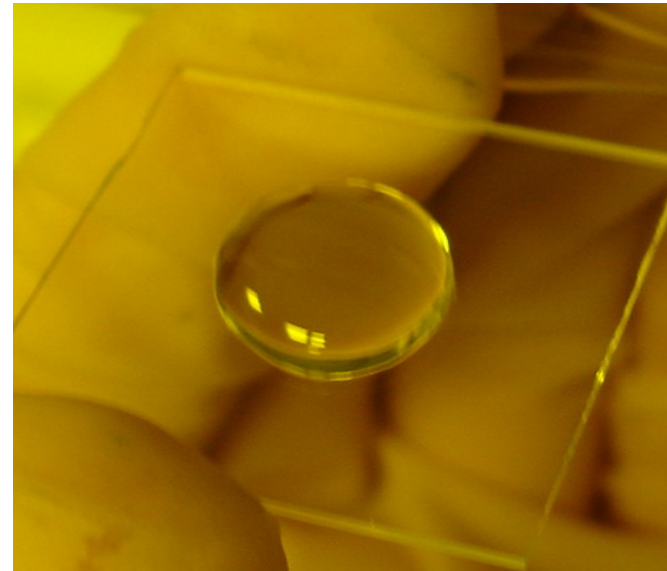
Solvent Compatibility of Poly(dimethylsiloxane)-Based Microfluidic Devices

Anal. Chem. 2003, 75, 6544–6554

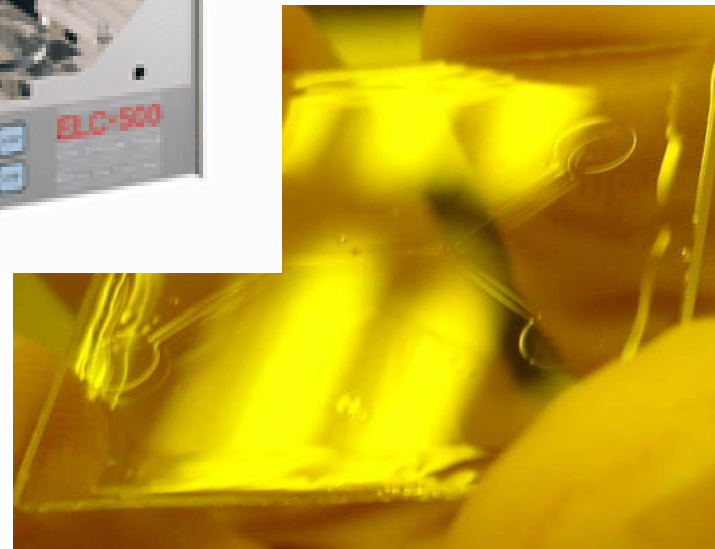
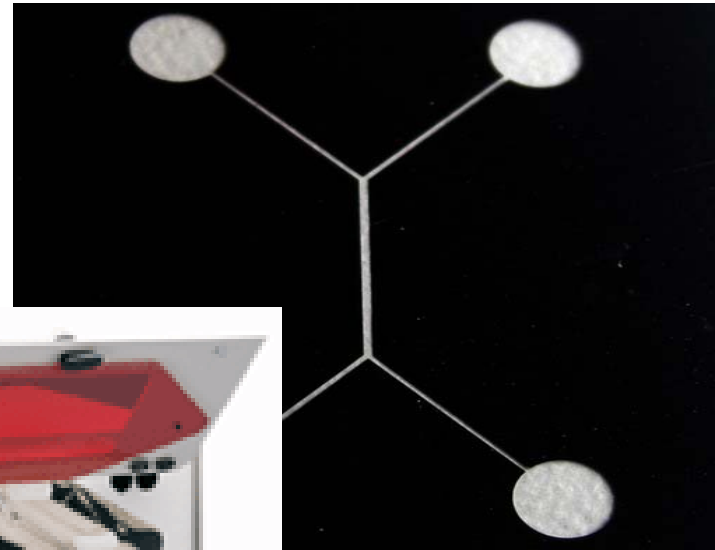
solvent	δ^a	S^b	μ (D)
perfluorotributylamine	5.6	1.00	0.0
perfluorodecalin	6.6	1.00	0.0
pentane	7.1	1.44	0.0
poly(dimethylsiloxane)	7.3	∞	0.6–0.9
diisopropylamine	7.3	2.13	1.2
hexanes	7.3	1.35	0.0
<i>n</i> -heptane	7.4	1.34	0.0
triethylamine	7.5	1.58	0.7
ether	7.5	1.38	1.1
cyclohexane	8.2	1.33	0.0
trichloroethylene	9.2	1.34	0.9
dimethoxyethane (DME)	8.8	1.32	1.6
xylenes	8.9	1.41	0.3
toluene	8.9	1.31	0.4
ethyl acetate	9.0	1.18	1.8
benzene	9.2	1.28	0.0
chloroform	9.2	1.39	1.0
2-butanone	9.3	1.21	2.8
tetrahydrofuran (THF)	9.3	1.38	1.7
dimethyl carbonate	9.5	1.03	0.9
chlorobenzene	9.5	1.22	1.7
methylene chloride	9.9	1.22	1.6
acetone	9.9	1.06	2.9
dioxane	10.0	1.16	0.5
pyridine	10.6	1.06	2.2
<i>N</i> -methylpyrrolidone (NMP)	11.1	1.03	3.8
<i>tert</i> -butyl alcohol	10.6	1.21	1.6
acetonitrile	11.9	1.01	4.0
1-propanol	11.9	1.09	1.6
phenol	12.0	1.01	1.2
dimethylformamide (DMF)	12.1	1.02	3.8
nitromethane	12.6	1.00	3.5
ethyl alcohol	12.7	1.04	1.7
dimethyl sulfoxide (DMSO)	13.0	1.00	4.0
propylene carbonate	13.3	1.01	4.8
methanol	14.5	1.02	1.7
ethylene glycol	14.6	1.00	2.3
glycerol	21.1	1.00	2.6
water	23.4	1.00	1.9



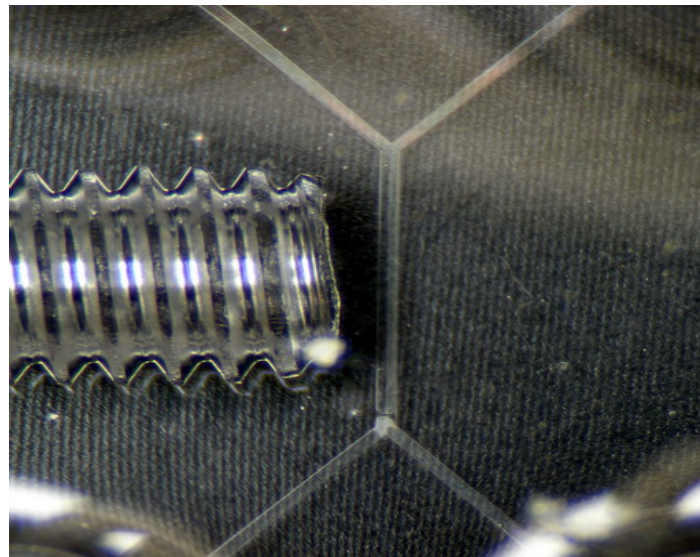
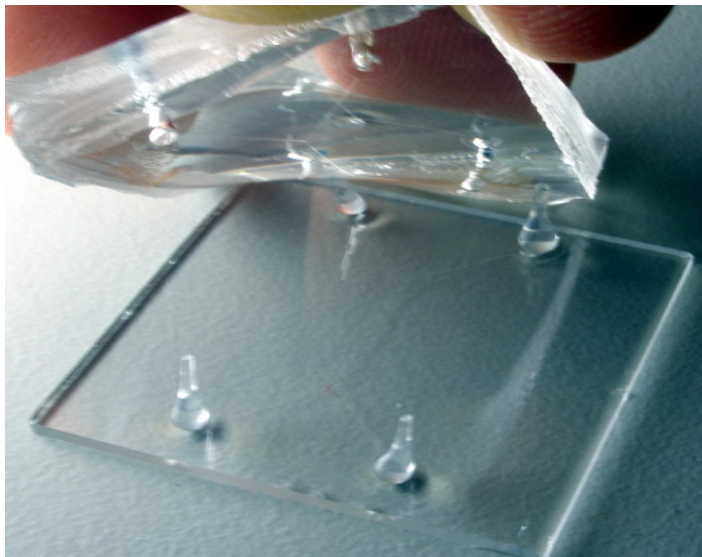
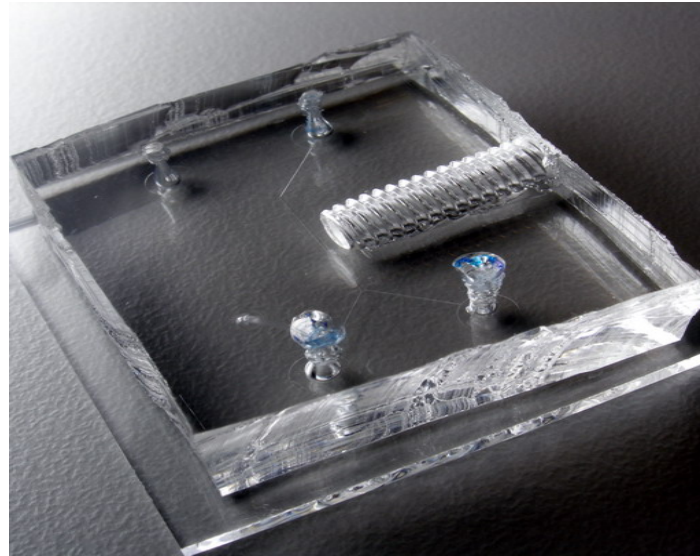
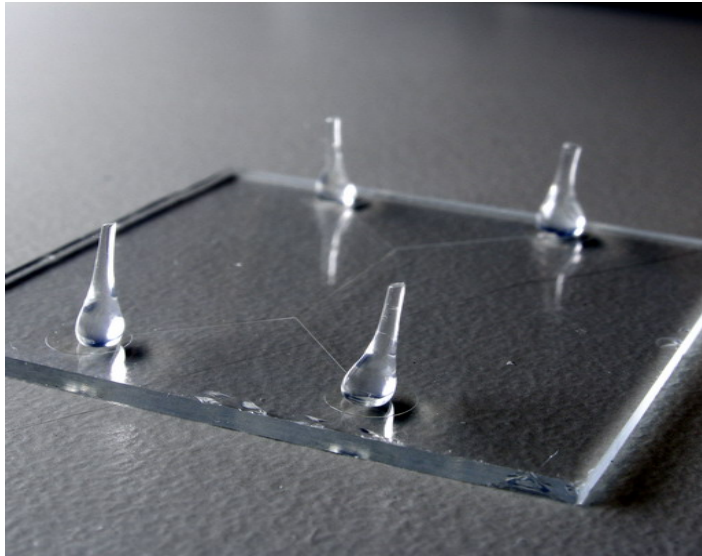
Microfluidics by Soft Lithography



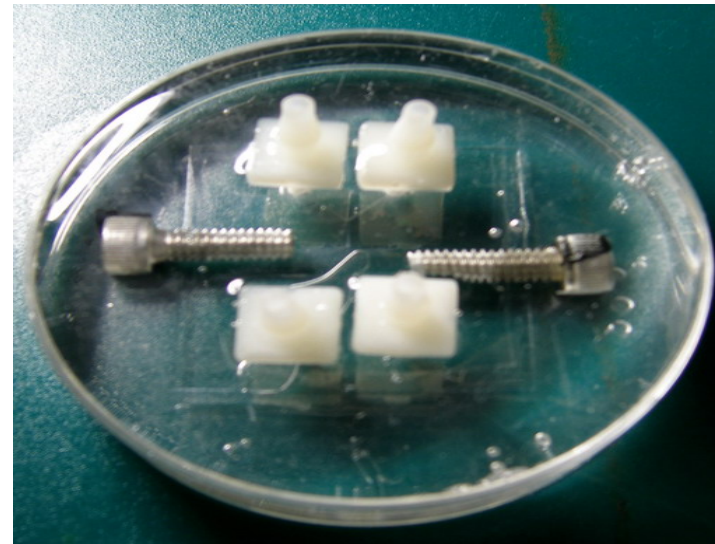
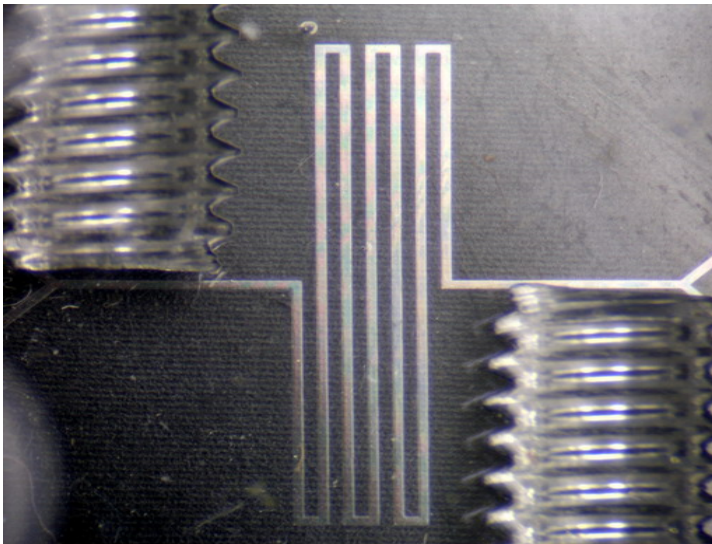
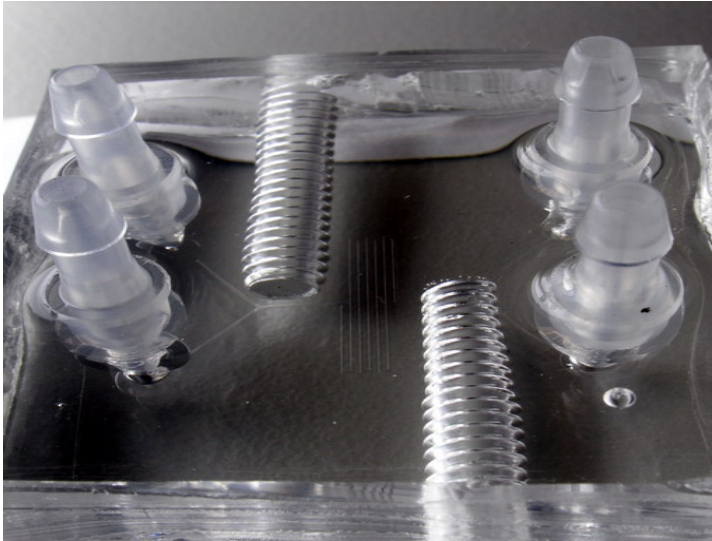
Microfluidics by Soft Lithography



Microfluidics by Soft Lithography

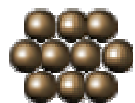
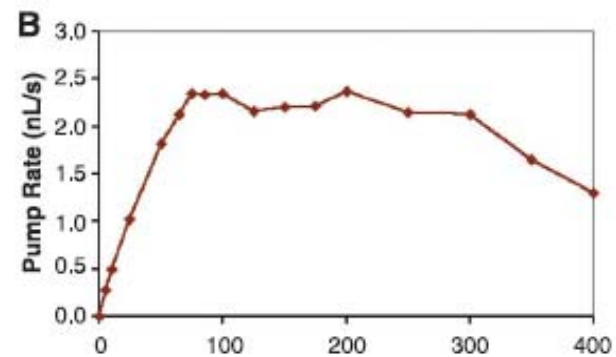
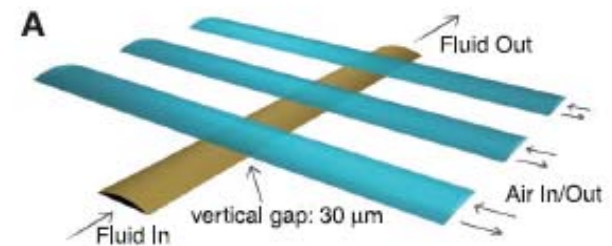
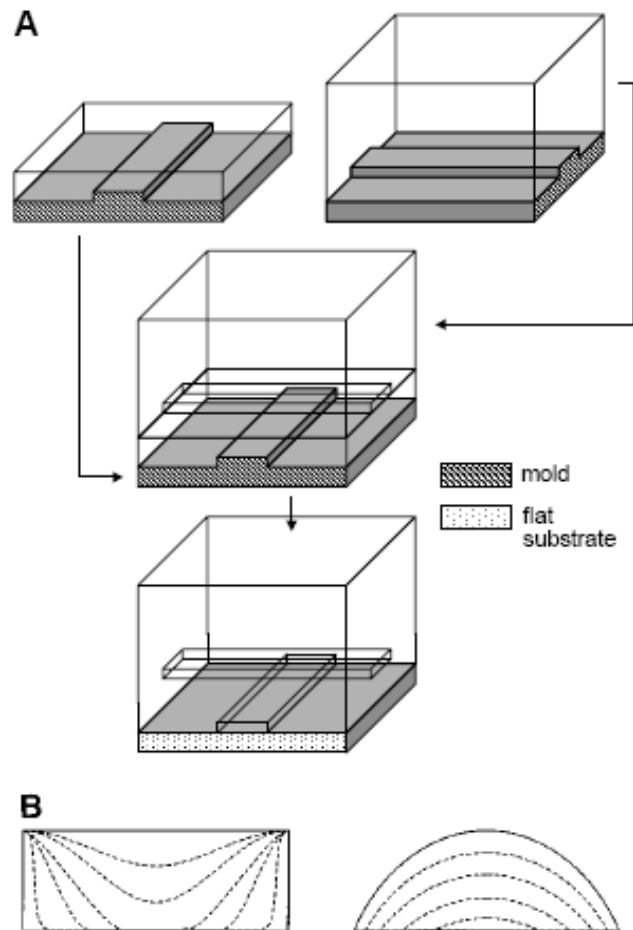


Microfluidics by Soft Lithography



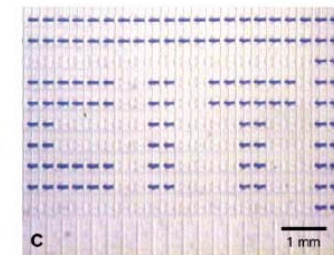
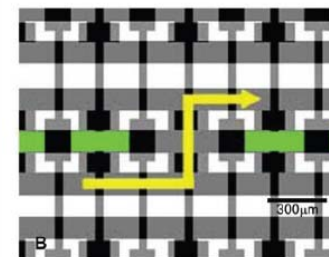
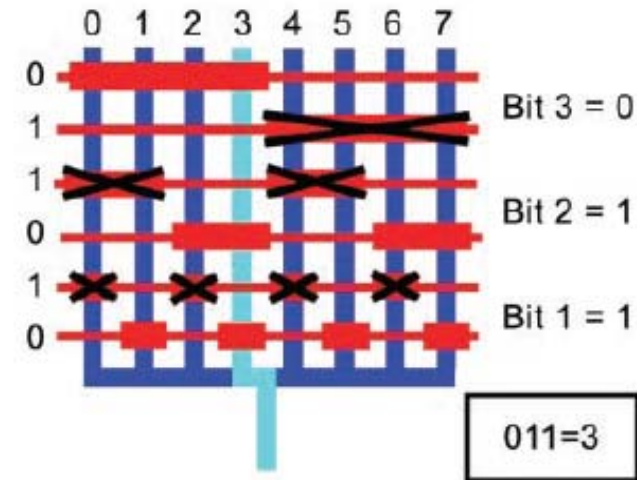
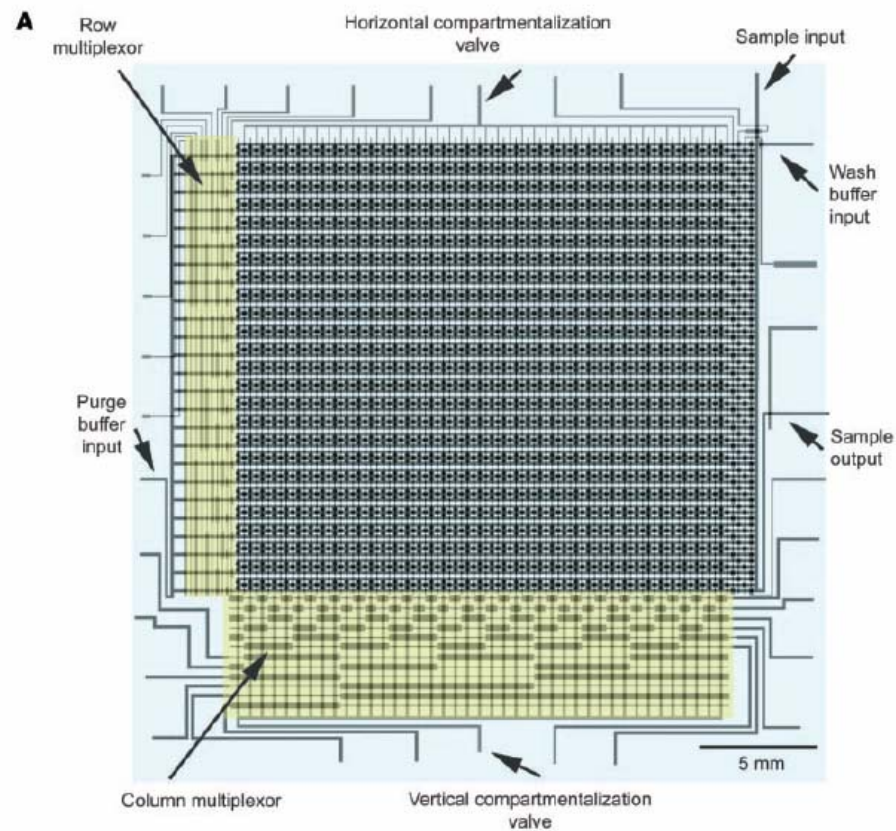
Monolithic Microfabricated Valves and Pumps by Multilayer Soft Lithography

SCIENCE VOL 288 7 APRIL 2000



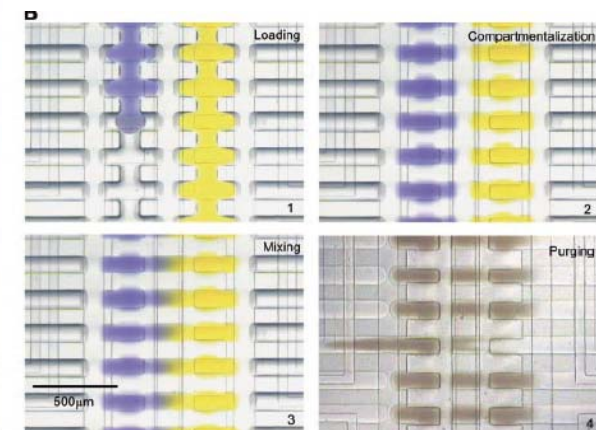
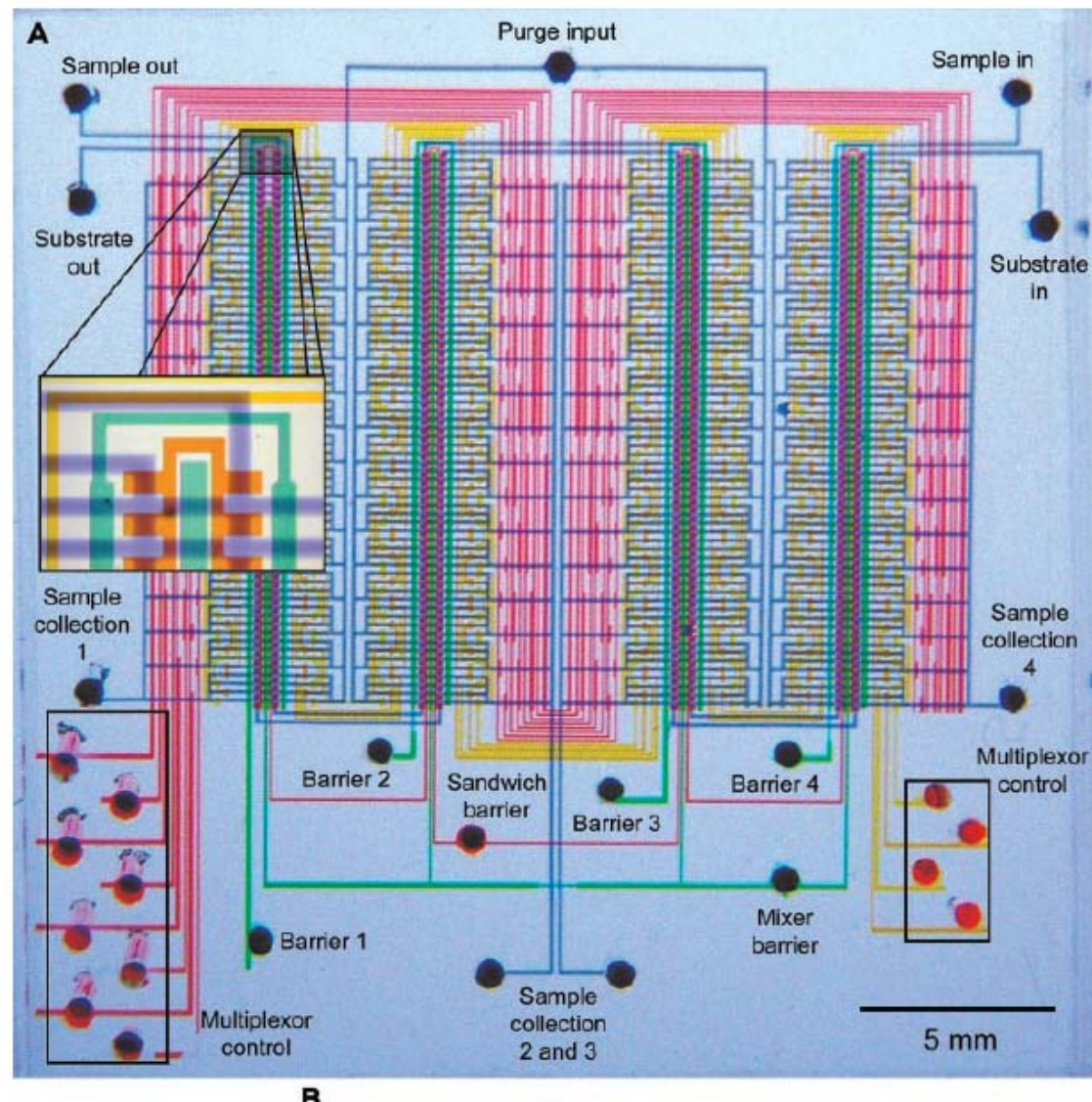
Microfluidic Large-Scale Integration

18 OCTOBER 2002 VOL 298 SCIENCE



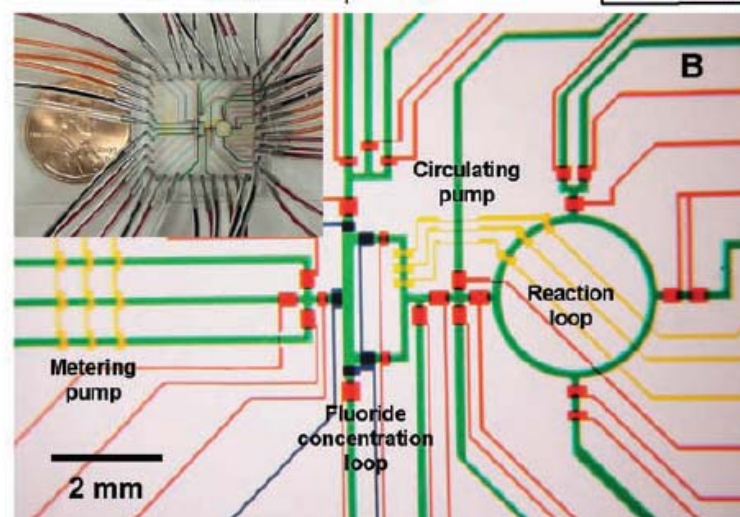
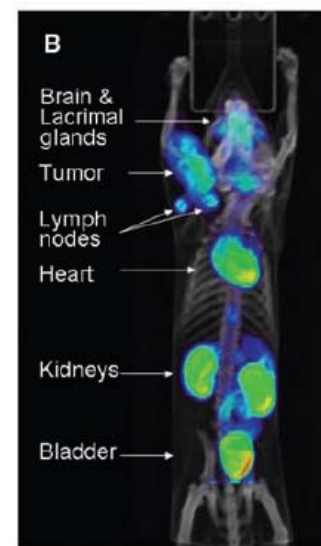
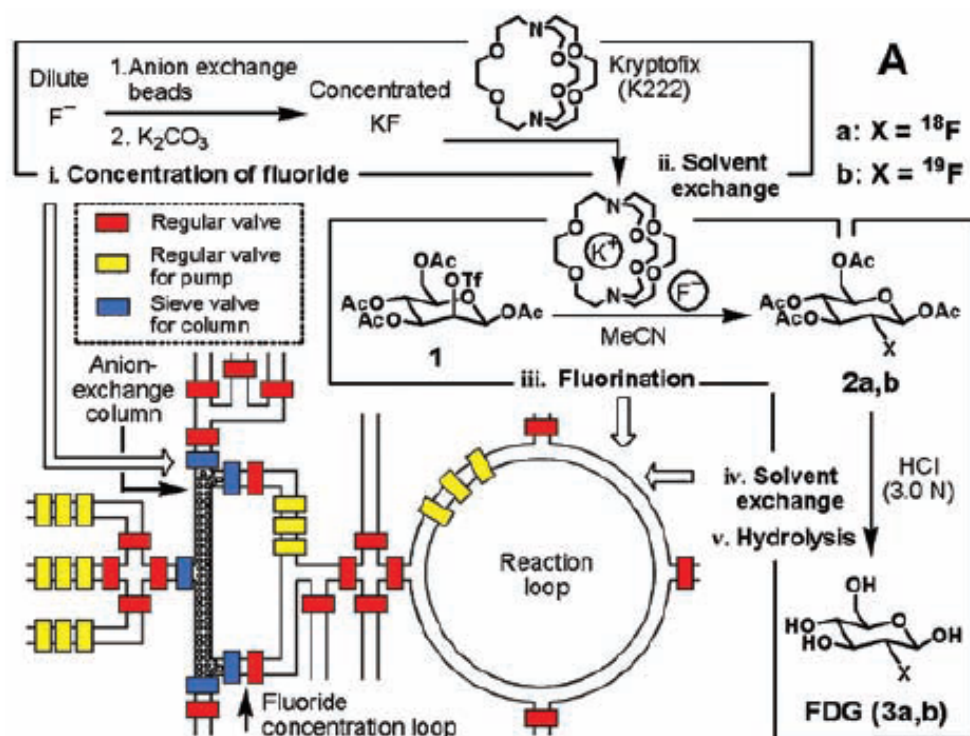
$2 \log_2 n$ control channels.



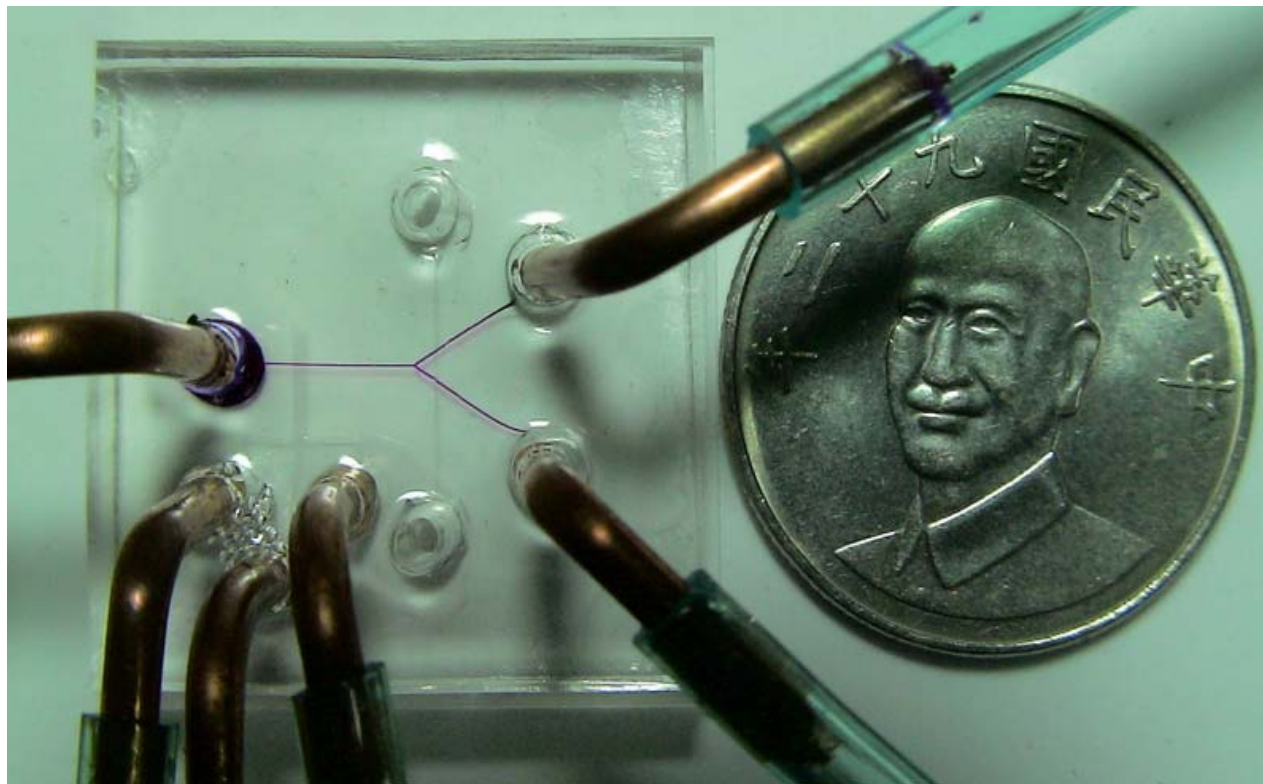


Multistep Synthesis of a Radiolabeled Imaging Probe Using Integrated Microfluidics

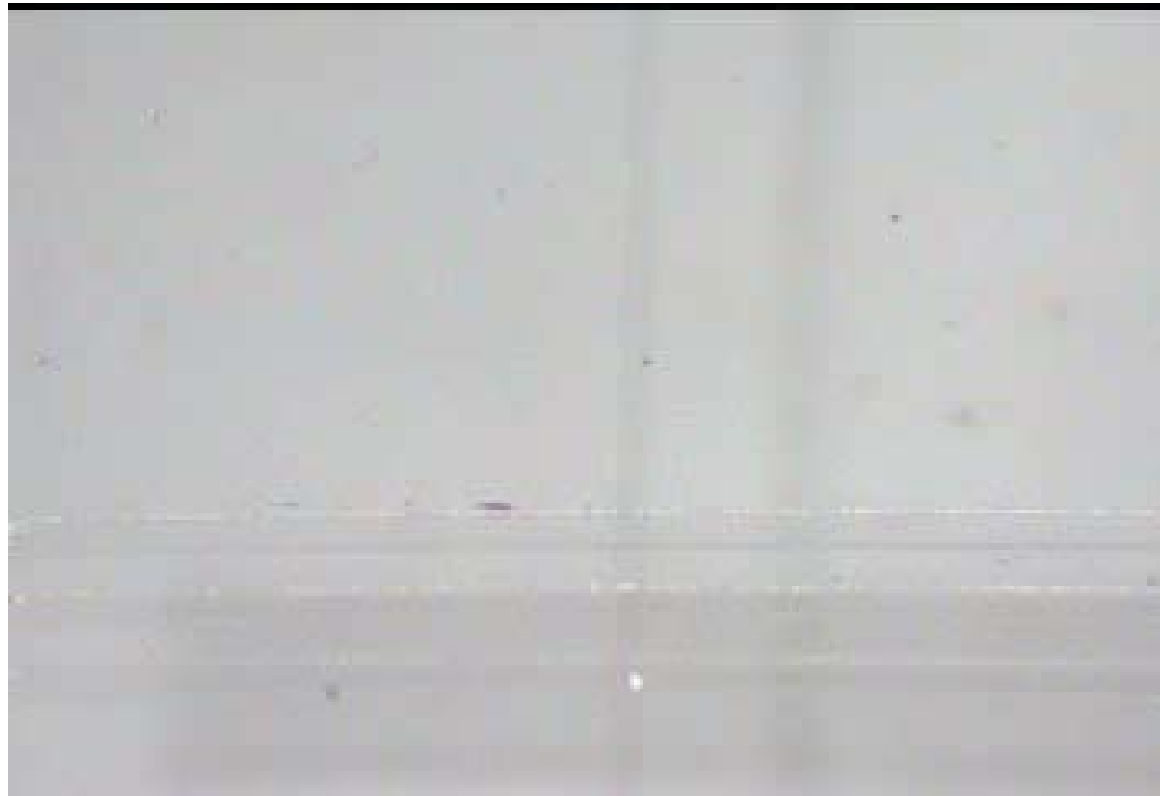
SCIENCE VOL 310 16 DECEMBER 2005



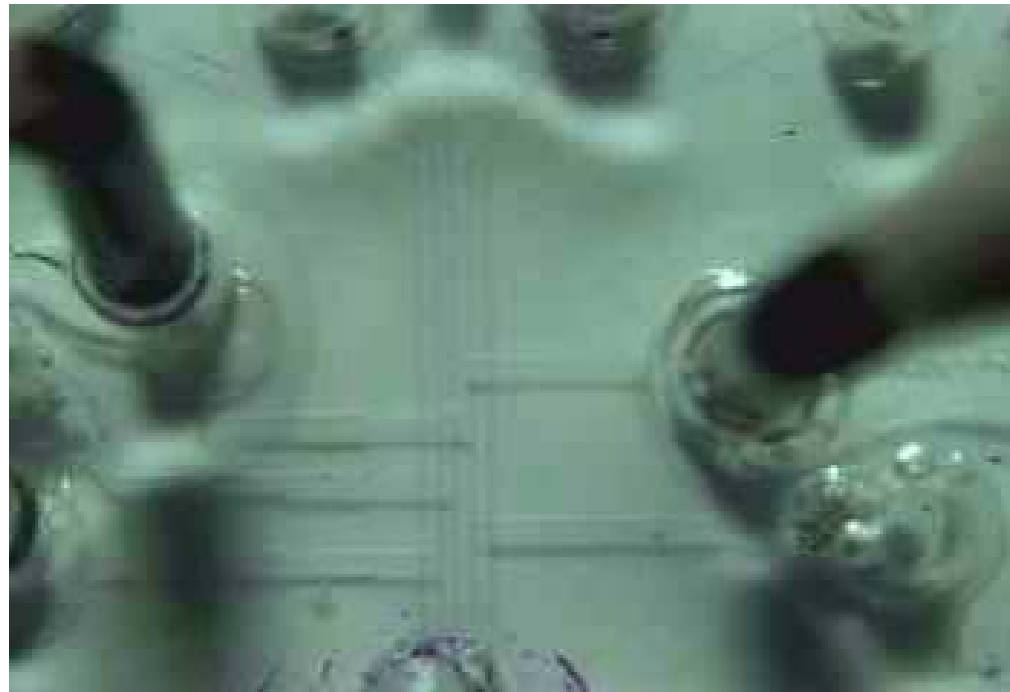
Microfabricated Fluidic System by Soft Lithography



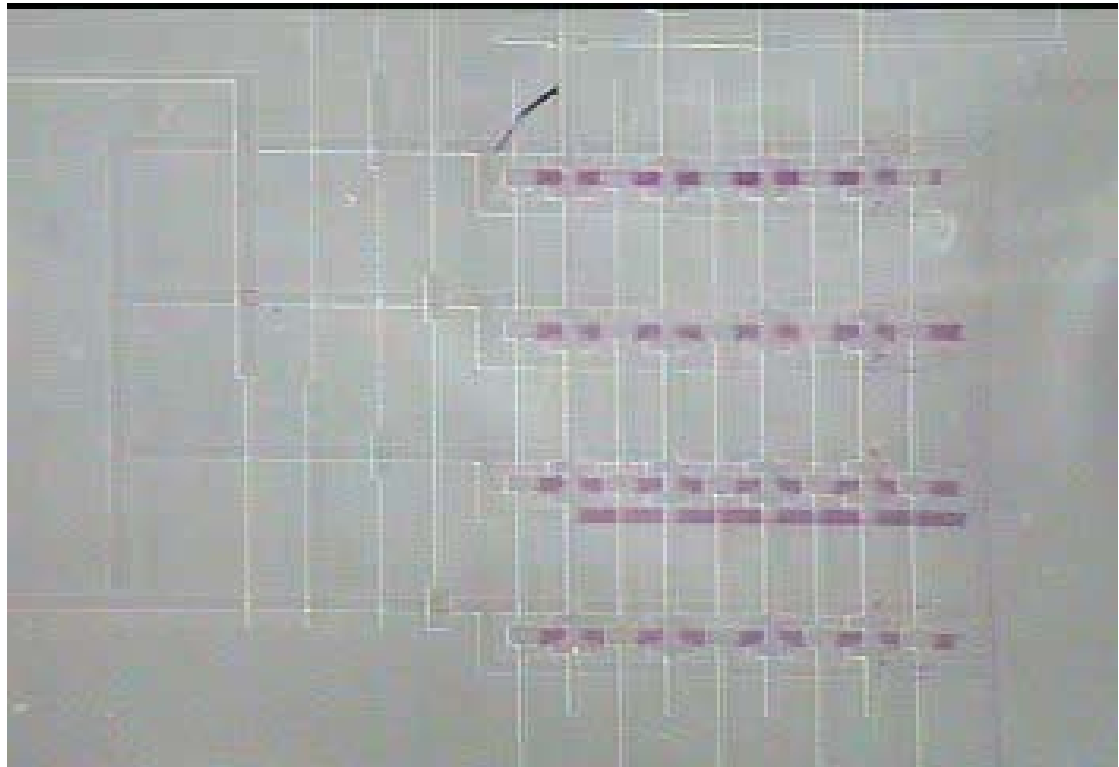
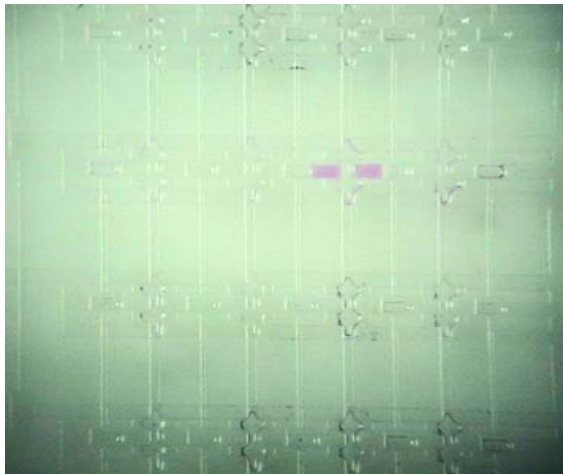
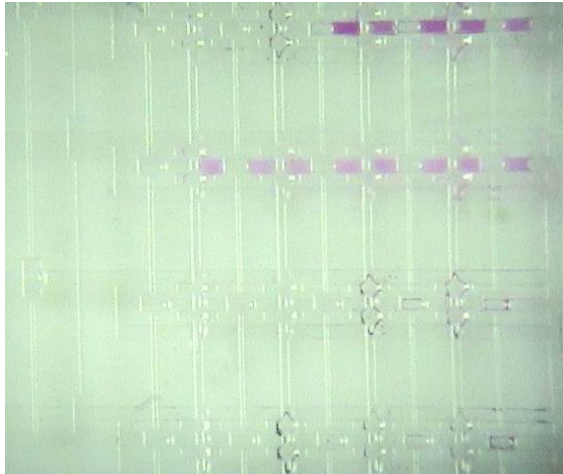
Peristaltic Pump by Soft Lithography



Fluidic Control by Microfabricated Valves



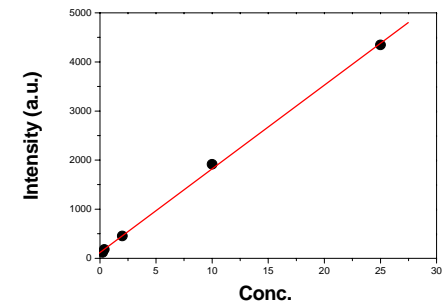
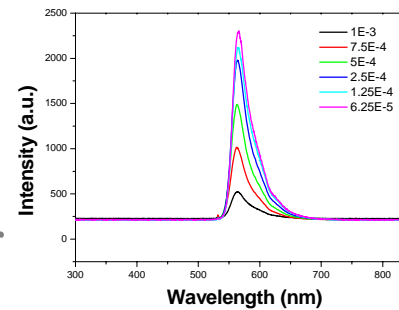
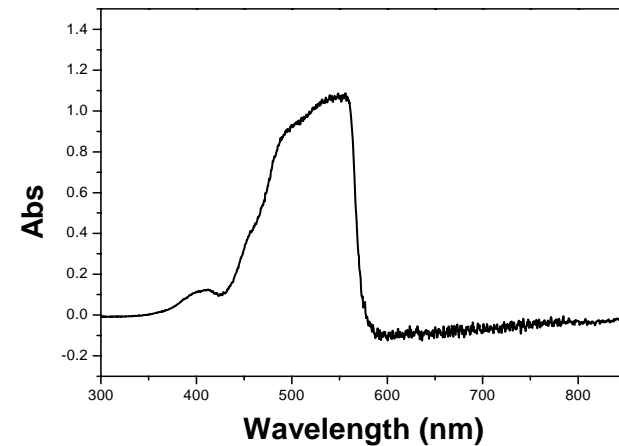
Addressable Microfluidic System



Integrated Detection System in Microchannel



fluidic channel + microlens + fiber



Reference

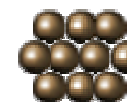
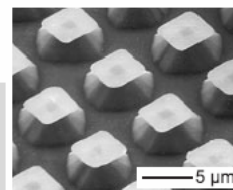
Adv. Mater. 2004, 16, No. 15, August 4 1249

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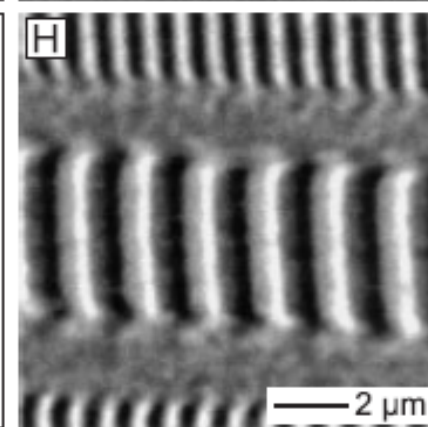
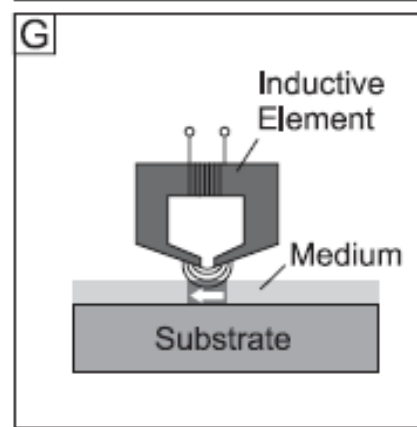
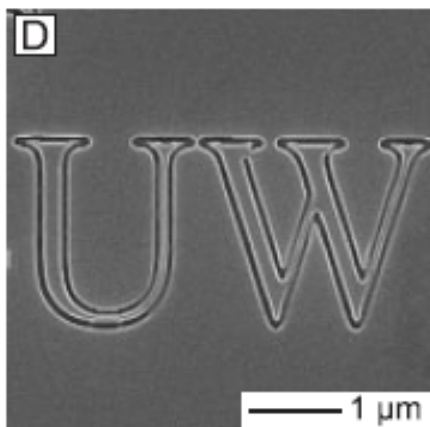
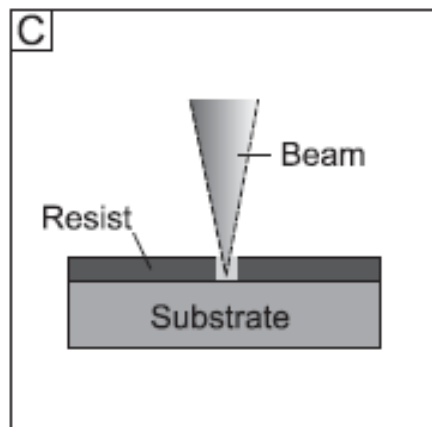
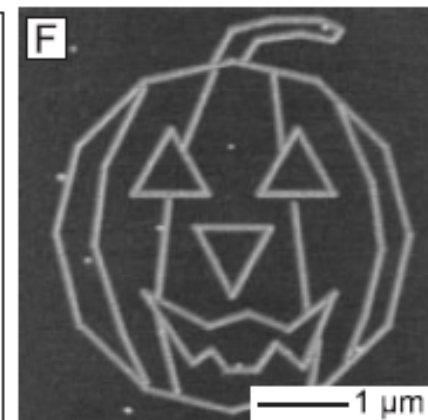
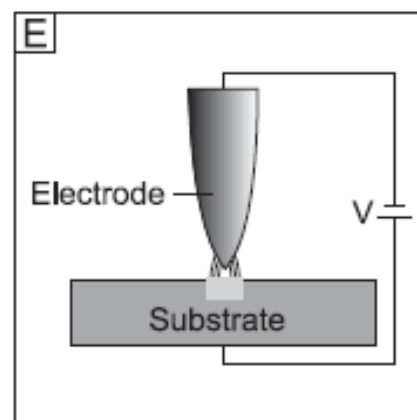
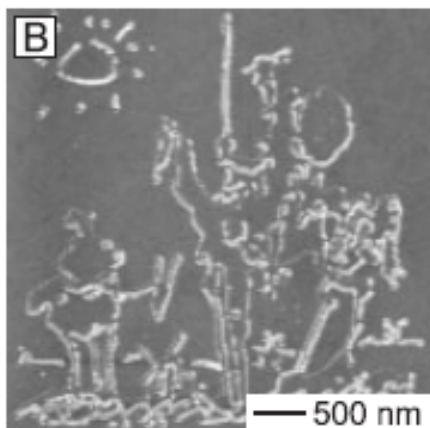
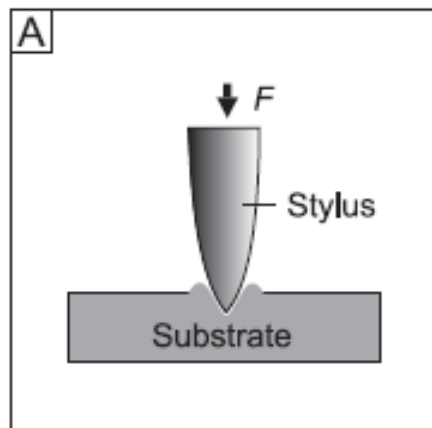
Patterning: Principles and Some New Developments**

By Matthias Geissler and Younan Xia*

This article provides an overview of various patterning methodologies, and it is organized into three major sections: generation of patterns, replication of patterns, and three-dimensional patterning. Generation of patterns from scratch is usually accomplished by serial techniques that are able to provide arbitrary features. The writing process can be carried out in many different ways. It can be achieved using a rigid stylus; or a focused beam of photons, electrons, and other energetic particles. It can also be accomplished using an electrical or magnetic field; or through localized add-on of materials such as a liquid-like ink from an external source. In addition, some ordered but relatively simple patterns can be formed by means of self-assembly. In replication of patterns, structural information from a mask, master, or stamp is transferred to multiple copies with the use of an appropriate material. The patterned features on a mask are mainly used to direct a flux of radiation or physical matter from a source onto a substrate, whereas a master/stamp serves as the original for replication based on embossing, molding, or printing. The last section of this article deals with three-dimensional patterning, where both vertical and lateral dimensions of a structure need to be precisely controlled to generate well-defined shapes and profiles. The article is illustrated with various examples derived from recent developments in this field.

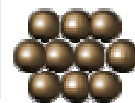


Direct Writing

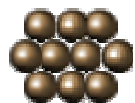
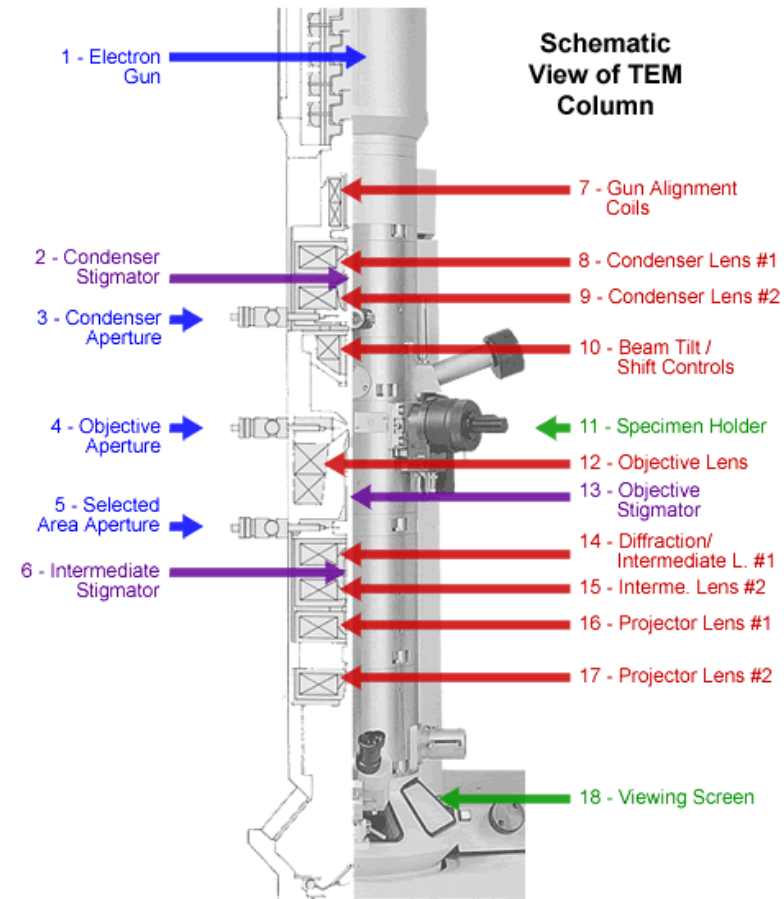




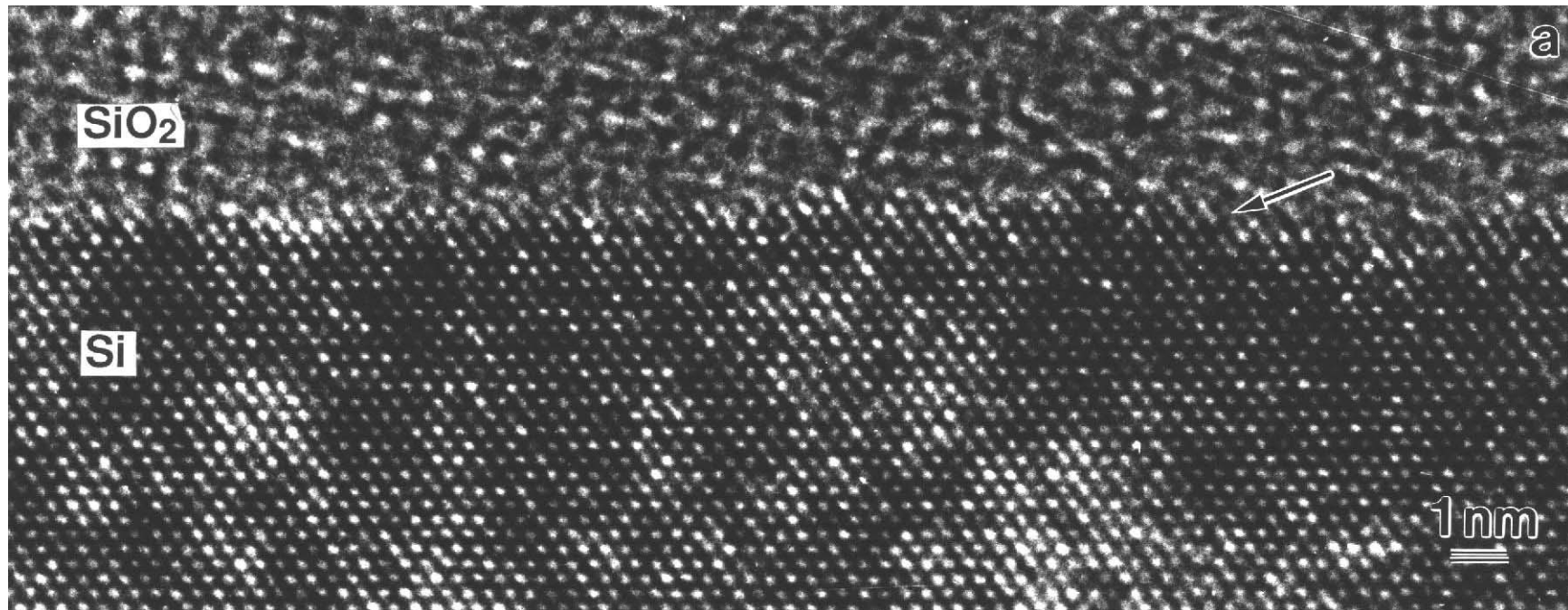
	write head				
	2mm	4mm	10mm	20mm	40mm
min. structure size [μm]	0.6	1.0	2.5	5.0	10.0
address resolution [nm]	20	40	100	200	400
writing speed [mm^2/min]	1.5	5.7	36	119	416
line width uniformity, 3σ [nm]	80	100	220	440	880
edge roughness, 3σ [nm]	60	80	120	180	280



Electron Microscope



TEM Image

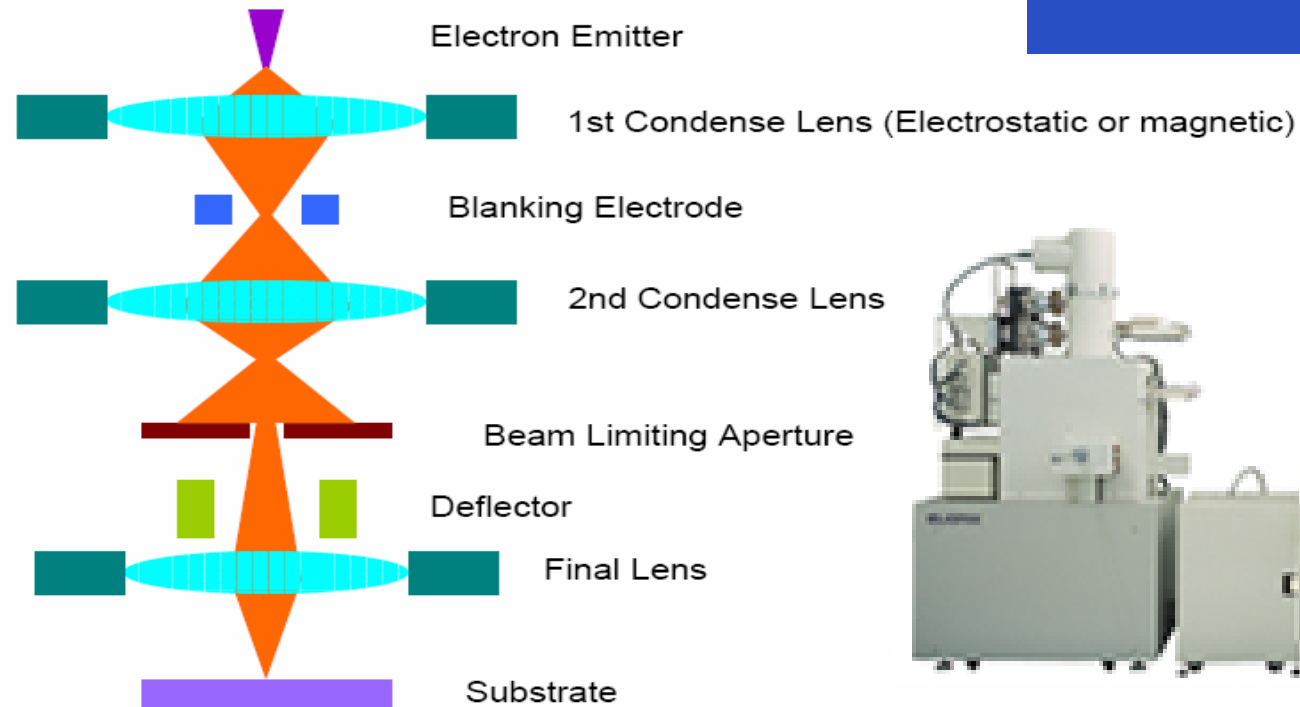


E-Beam Lithography

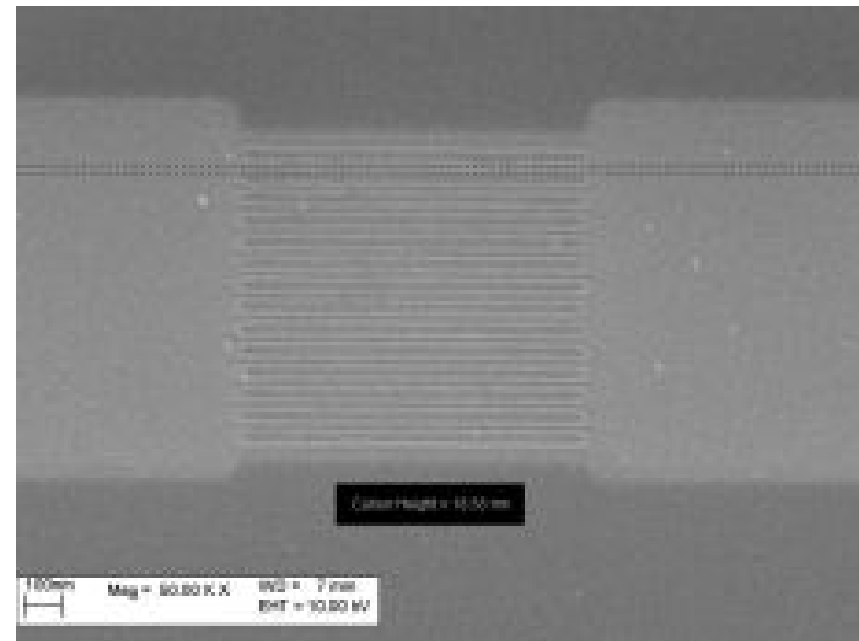
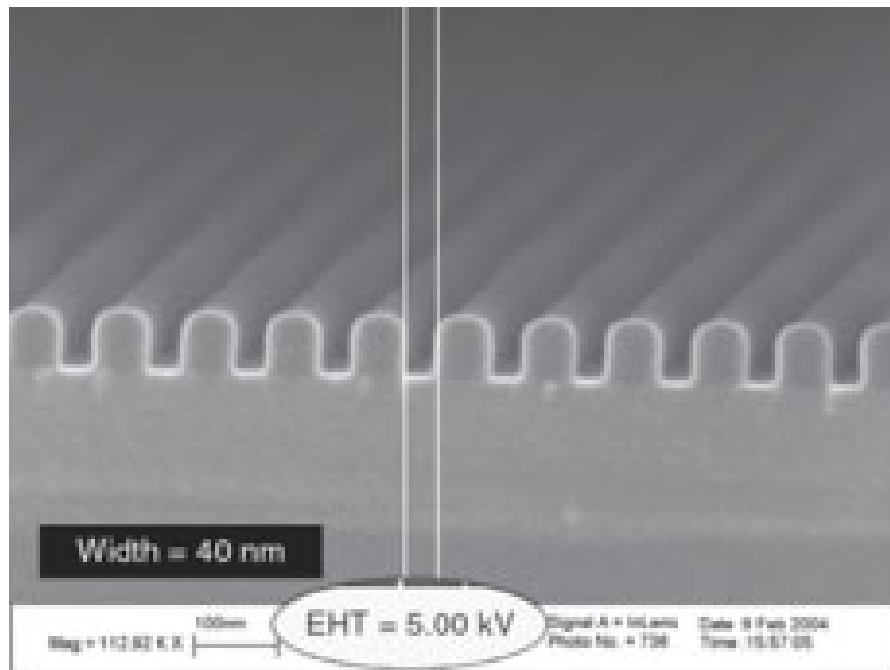
Resolution:

2 nm @ 20 KeV

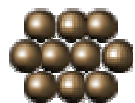
2 nm @ 1.0 KeV



E-beam Writer

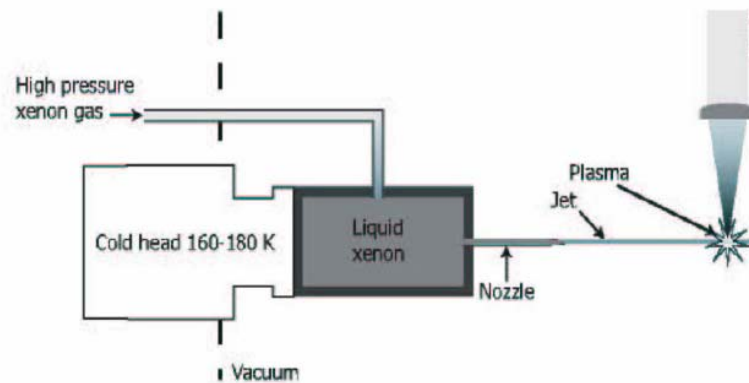


Better than 10 nm lines over 4 inch wafer



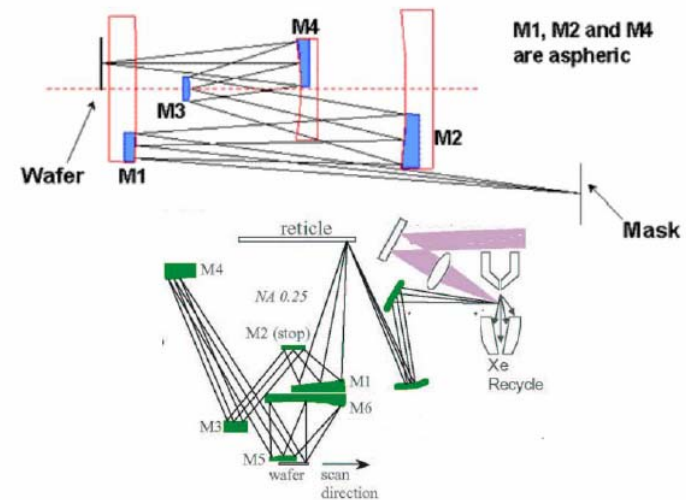
EUV System

EUV Source

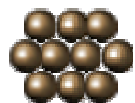


1. 13.4nm source.
2. Laser-produced plasma

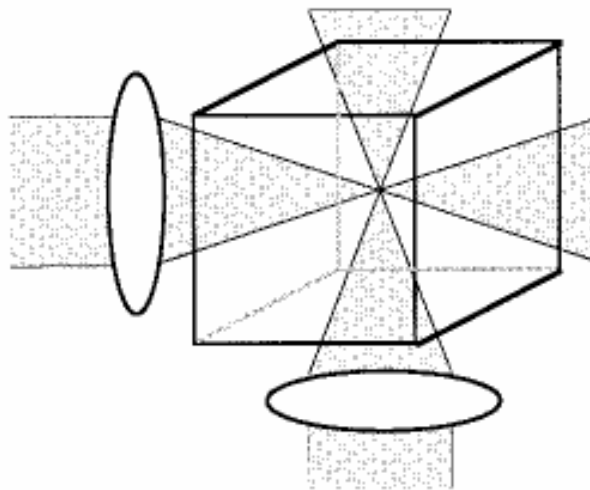
Reflective Optics for EUV



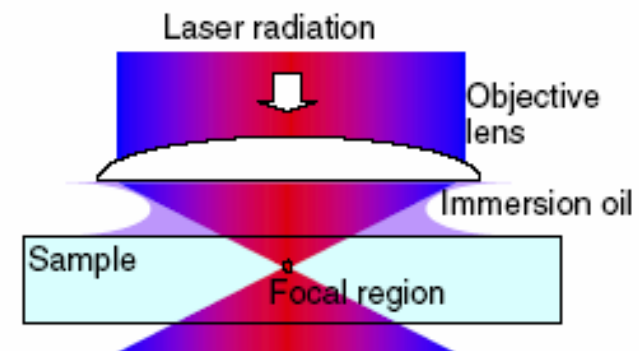
EUV light ($< 100\text{nm}$) is absorbed by almost all materials \rightarrow No lens.



Two Photon Writing

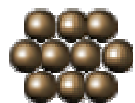


(a)



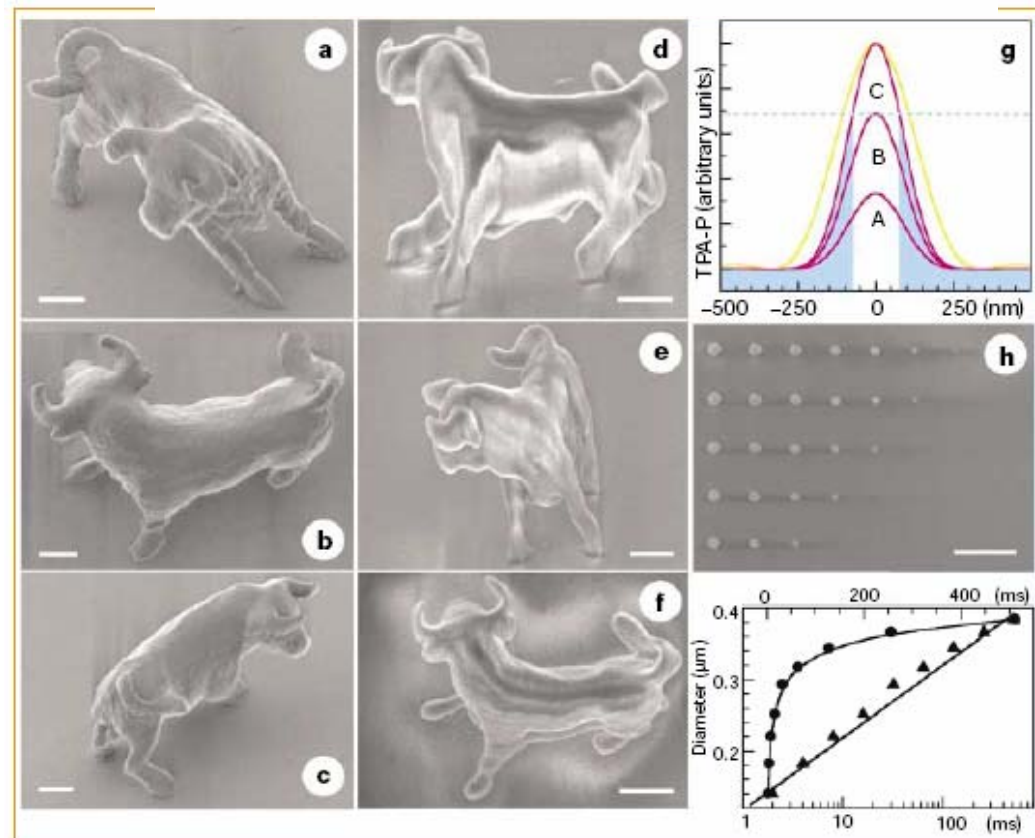
$$D(r) = w_0 \sqrt{2 \ln \left(\frac{I(r)}{I_{th}} \right) \frac{1}{N}},$$

$$L(z) = 2z_R \sqrt{\left(\frac{I(z)}{I_{th}} \right)^{\frac{1}{n}} - 1},$$



Two Photon Writing

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Two Photon Writing

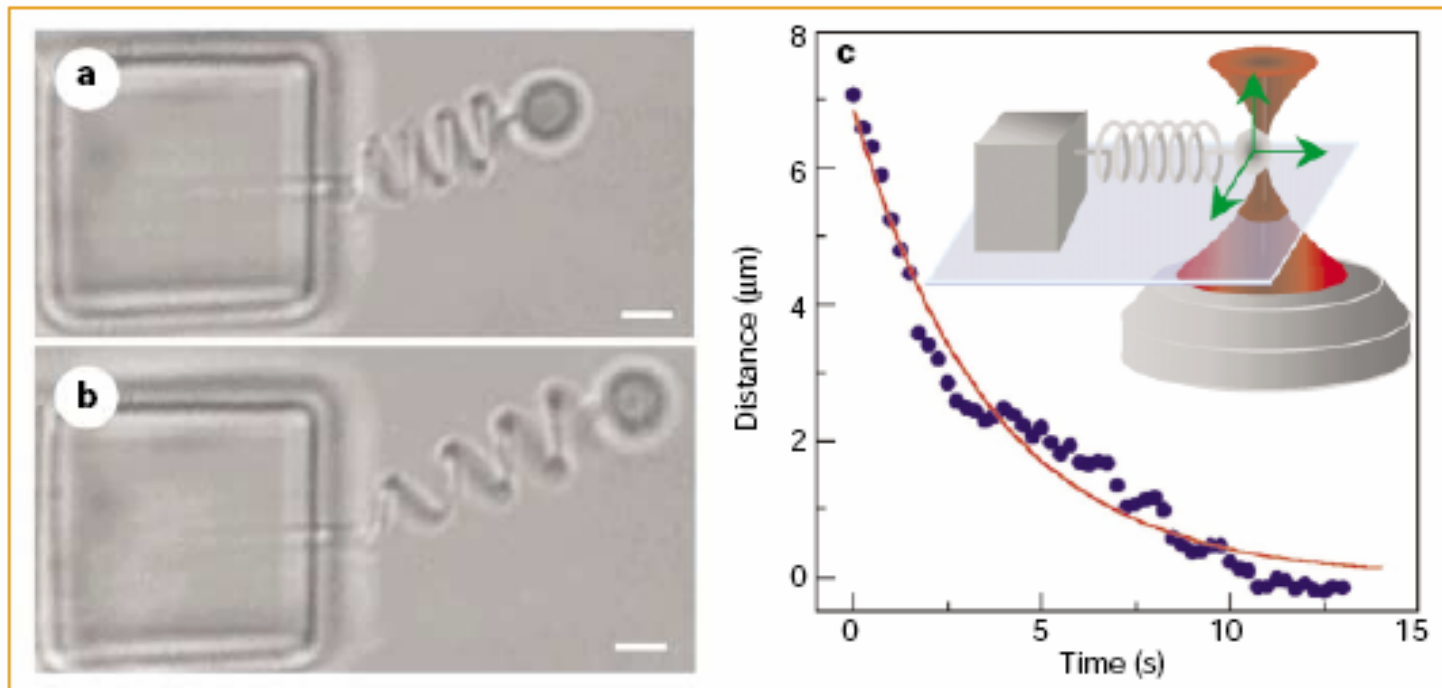
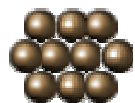


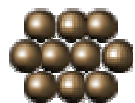
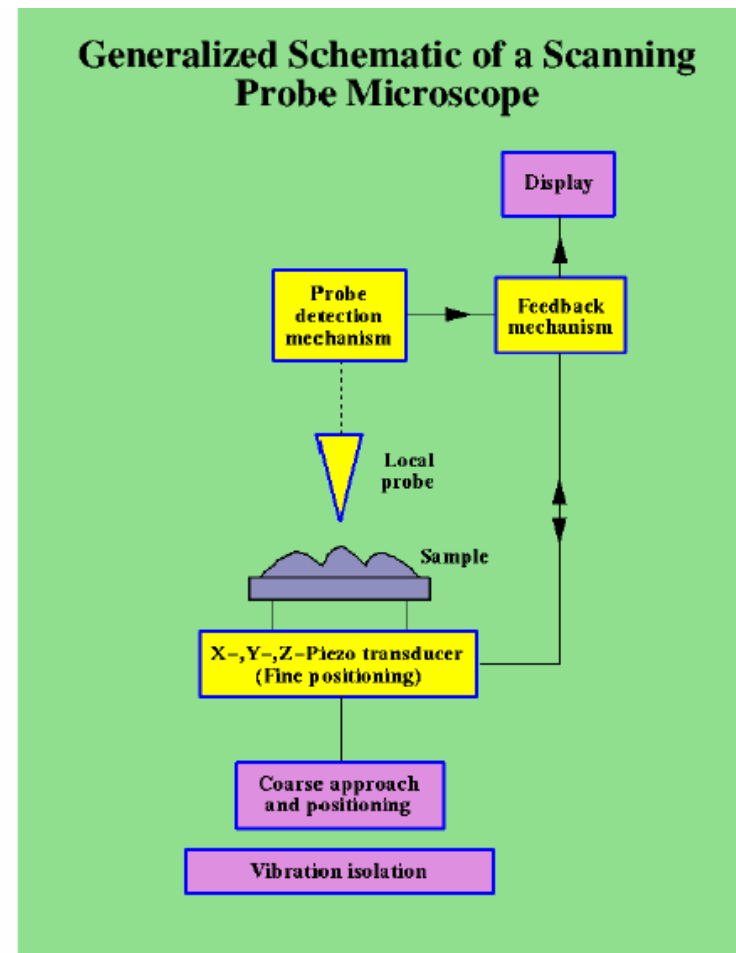
Figure 2 Functional micro-oscillator system, in which not only the spring but also the cubic anchor and the bead were produced using our two-photon absorption system. The oscillator was kept in ethanol so that the buoyancy would balance gravity and eliminate bead-substrate friction. **a, b**, The spring in its original (**a**) and extended (**b**) states. Scale bars, 2 μm . **c**, Restoring curve of the damping oscillation; inset, diagram showing driving of the oscillator by using laser trapping.



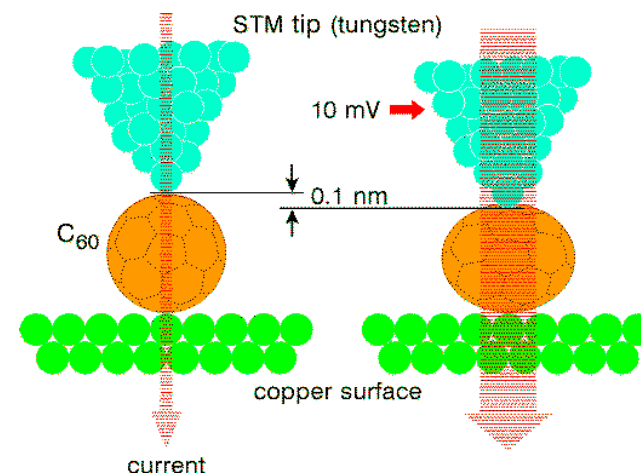
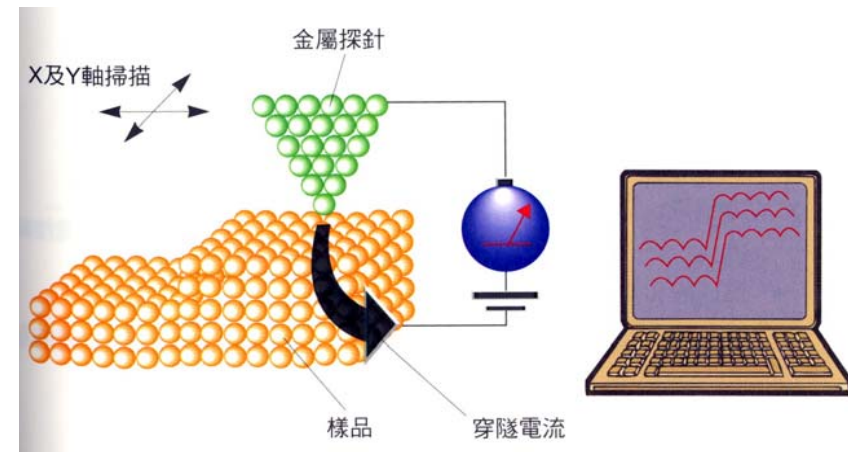
Introduction To Scanning Probe Microscopy

General features of SPM

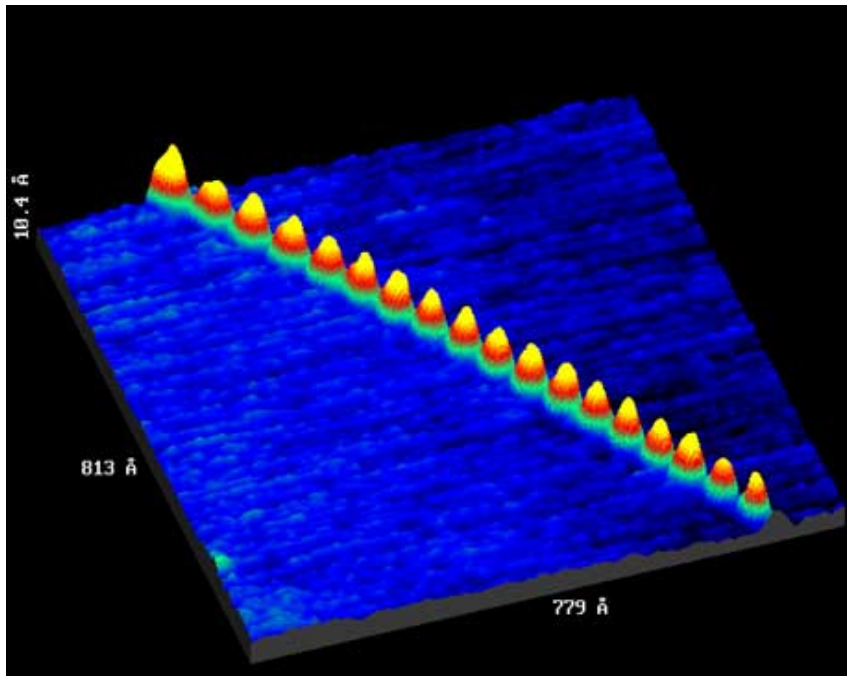
The first scanning probe microscope was invented in 1981 by Gerd Binnig and Heinrich Rohrer at the IBM laboratory in Zurich. Since that time, a vast family of scanning probe microscopes has been spawned, a few of which are represented in this family tree. Despite the huge number of types of SPM and modes in which they can be operated, the underlying operation is the same for them all. Each different type of SPM is characterized by the nature of the local probe and its interaction with the sample surface.



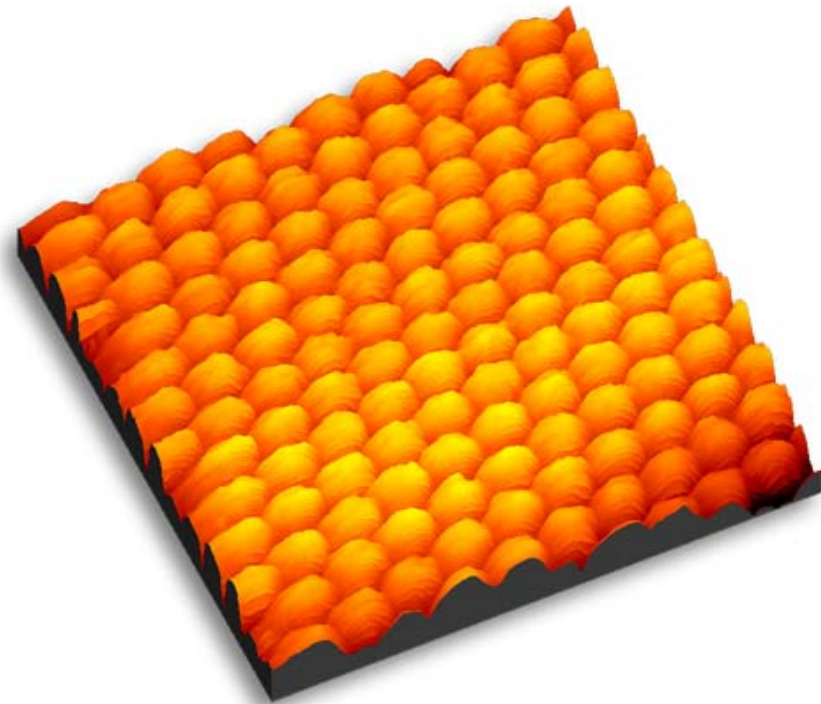
Scanning Tunneling Microscopy



STM Images



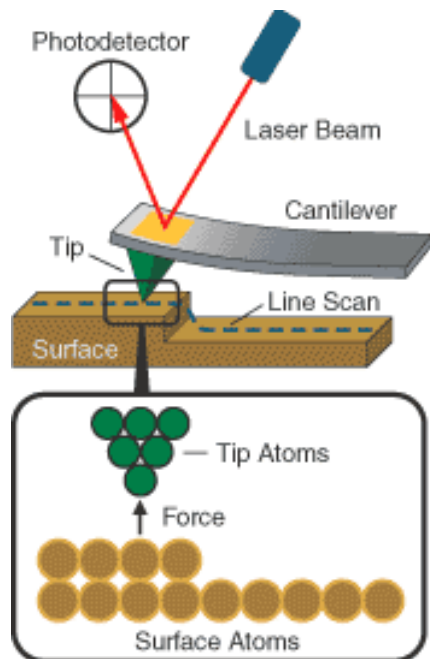
Polymer

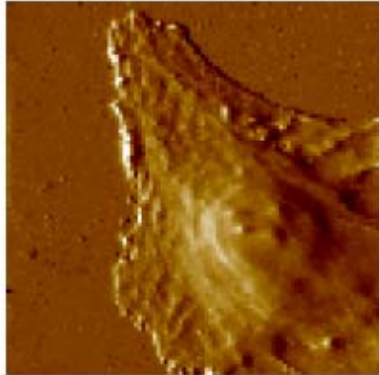


Gold atom



Atomic Force Microscopy





Cell surface imaging
AFM imaging of living
and fixed animal cells

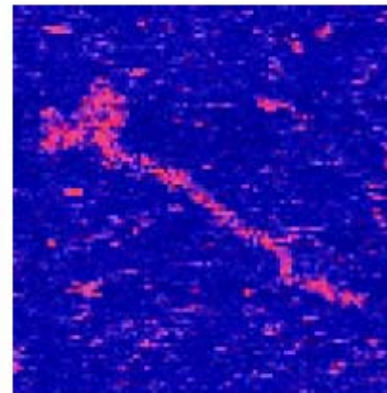


Chromosomes
A large selection of
work on both human
and plant
chromosomes

Scanning Probe Microscope



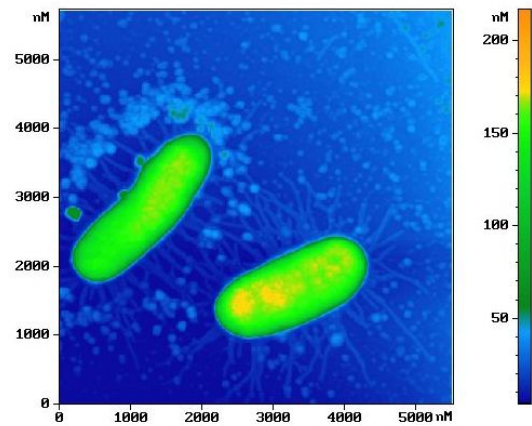
DNA
AFM images of
double-stranded
DNA and of some
special single-
double-stranded
DNA constructs



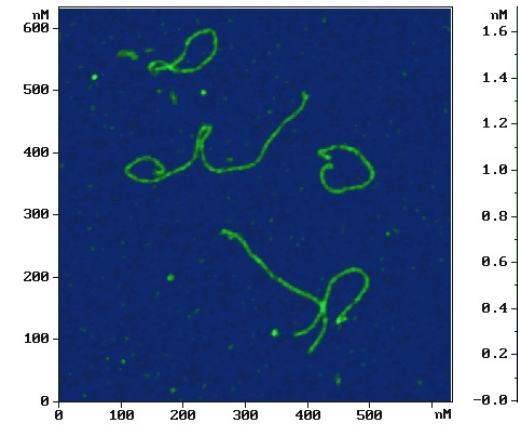
Muscle Proteins
High resolution
AFM images of
myosin and titin



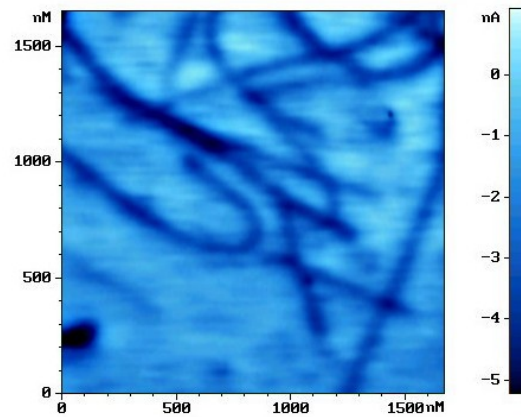
AFM Images



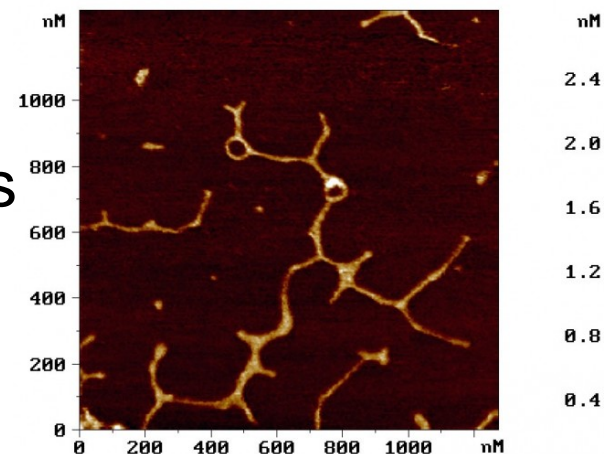
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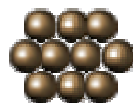
Protein



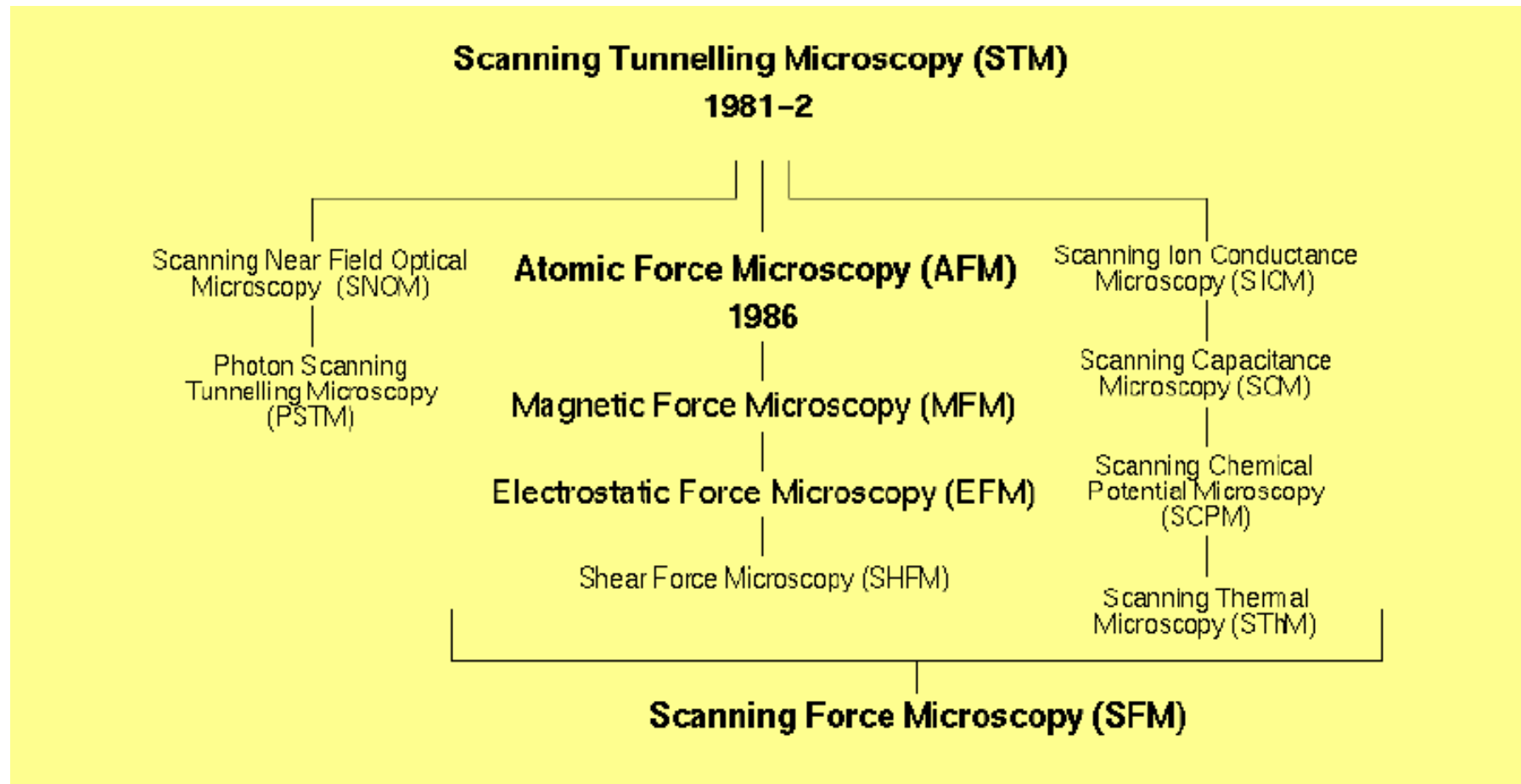
Nanotubes



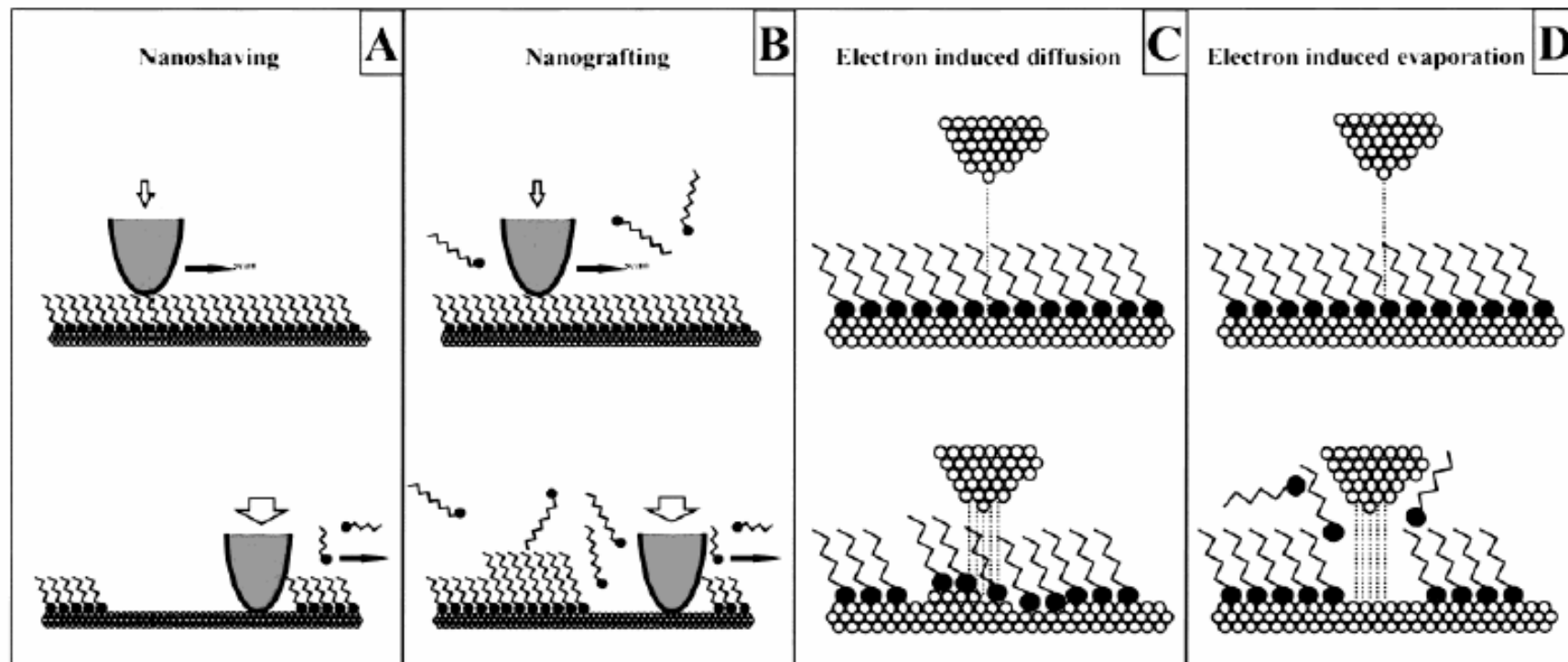
DNA



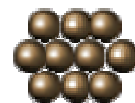
Scanning Probe Family



STM Lithography



Resist: Thiol



STM Lithography

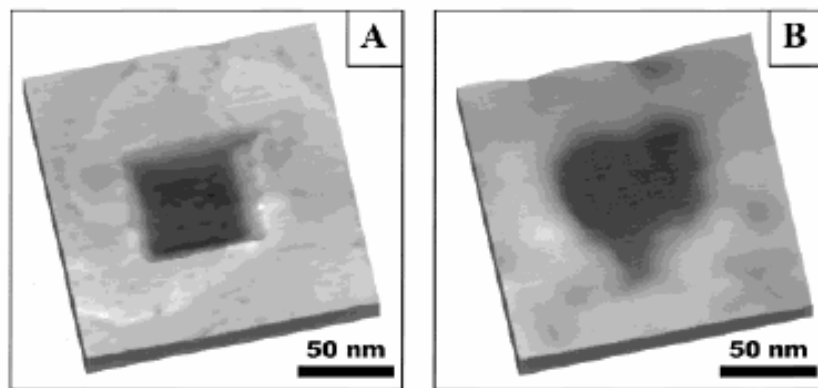
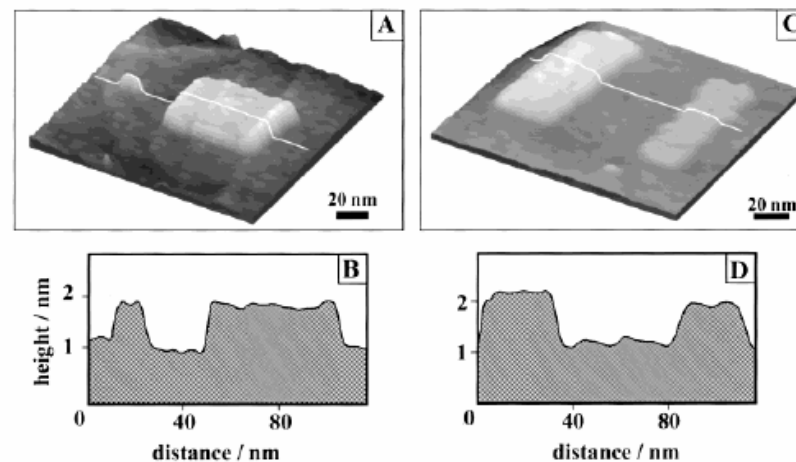


FIGURE 3. (A) $160 \times 160 \text{ nm}^2$ topographic images of $\text{C}_{18}\text{S}/\text{Au}(111)$ with the thiols shaved away from the central $50 \times 50 \text{ nm}^2$ square. (B) $160 \times 160 \text{ nm}^2$ topographic images of OTE/mica containing a heart-shaped pattern produced using nanoshaving.



Oxidation Lithography

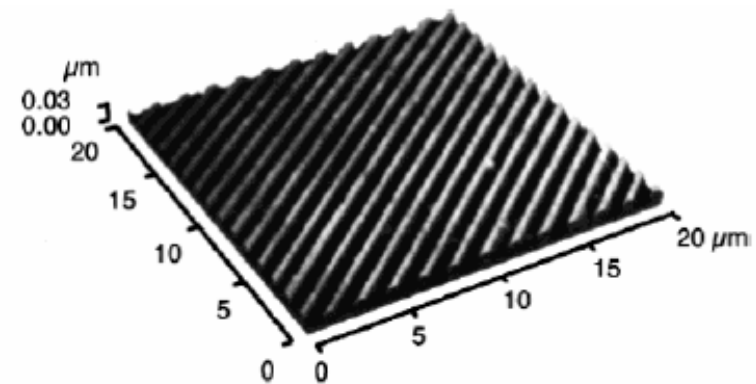
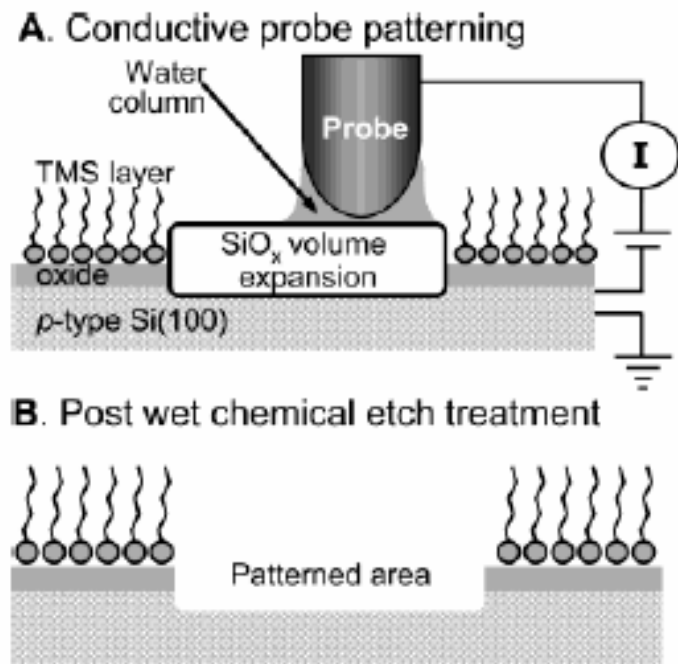
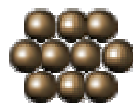


Figure 8. AFM images showing the results of AFM anodic oxidation and wet etching of a pattern in an ODS-SAM on a multilayer resist: scan speed, $10 \mu\text{m/s}$; probe current, 5 nA; etching steps with 0.5 wt % hydrofluoric acid (0.5 min) and 25 wt % tetraammonium hydroxide (3 min). Reprinted with permission from ref 111. Copyright 1999 Society of the American Institute of Physics.



AFM Lithography

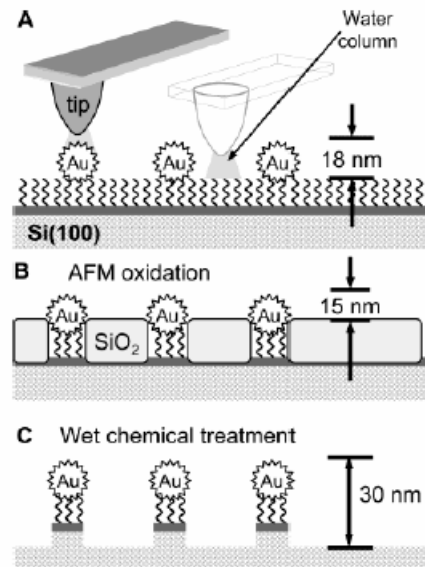


Figure 10. Schematic diagram of (A) selective anodization of the silicon regions not masked by nanoparticles. (B) The volume expansion of the silicon led to a decreased height contrast between the particles and the substrate. (C) After a wet etching step, silicon columns capped with a nanoparticle were formed.

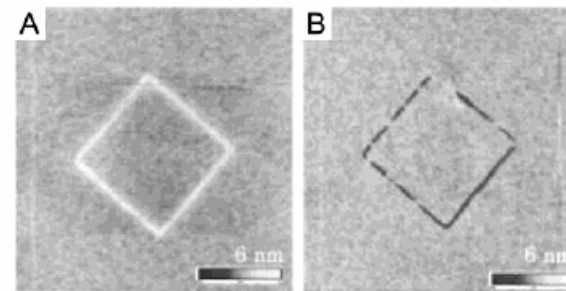
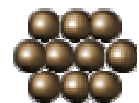


Figure 12. AFM images showing a lithographically defined pattern in a palmitic acid LB layer adsorbed to a SiO_x/Si substrate. Diamond pattern with protruding lines was written with -10 V applied tip bias (A), and that with grooves, with +10 V applied tip bias (B) (z-scale: 6 nm). Reprinted with permission from ref 123. Copyright 2002 American Institute of Physics.



Substitution Lithography

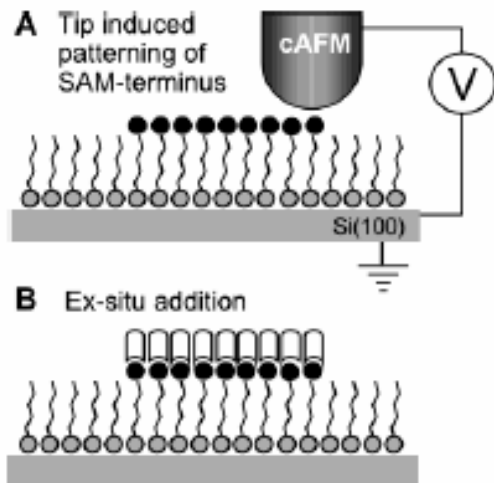
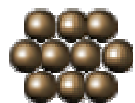
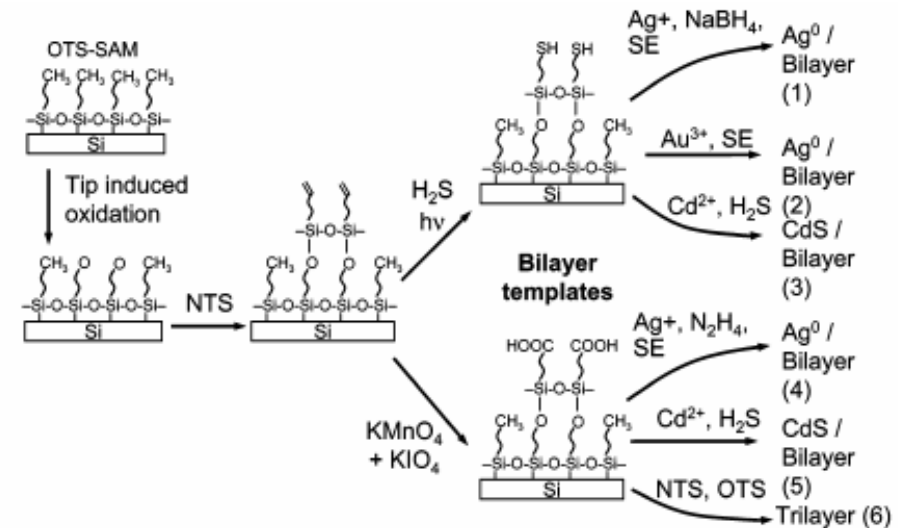
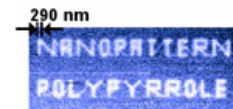
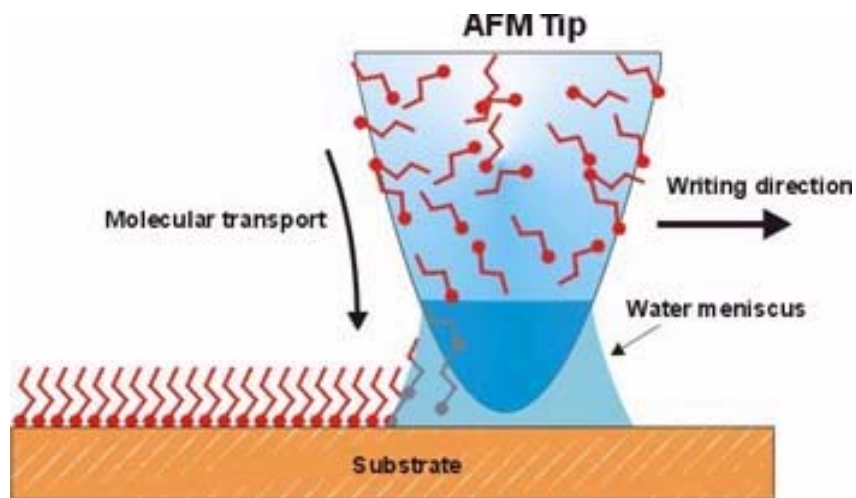


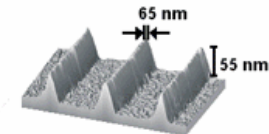
Figure 39. Schematic diagram of substitution lithography through the headgroup modification under an applied bias: (A) conductive AFM tip scanned under an applied bias voltage across a SAM and modified the headgroup without changing the structural integrity of the SAM; (B) altered functionality used for the subsequent binding of a second self-assembly molecule.



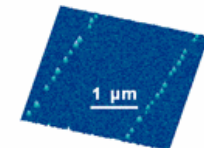
Dip-Pen Lithography



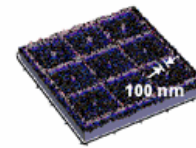
Conducting Polymers



Silicon Nanostructures

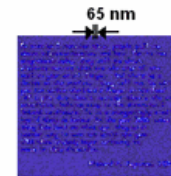
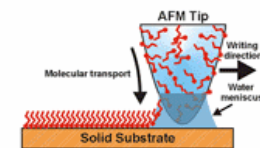


Single Nanoparticle Lines

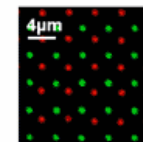


Protein Nanoarrays

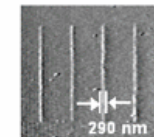
Dip-Pen Nanolithography



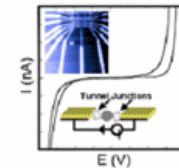
Small Organic Molecules



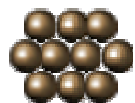
Ultrahigh Density DNA Arrays



Sol Gel Templates



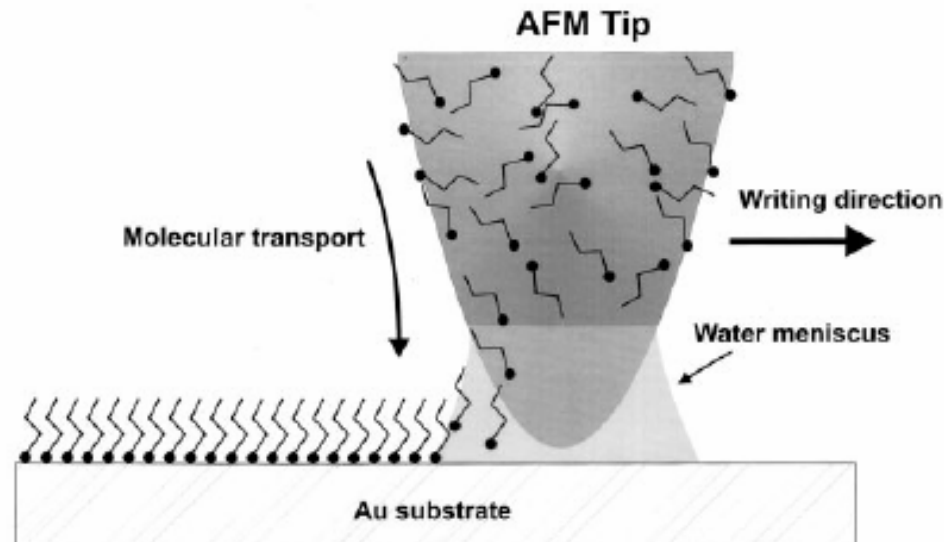
Single Particle Devices



"Dip-Pen" Nanolithography

Richard D. Piner, Jin Zhu, Feng Xu, Seunghun Hong,
Chad A. Mirkin*

Fig. 1. Schematic representation of DPN. A water meniscus forms between the AFM tip coated with ODT and the Au substrate. The size of the meniscus, which is controlled by relative humidity, affects the ODT transport rate, the effective tip-substrate contact area, and DPN resolution.

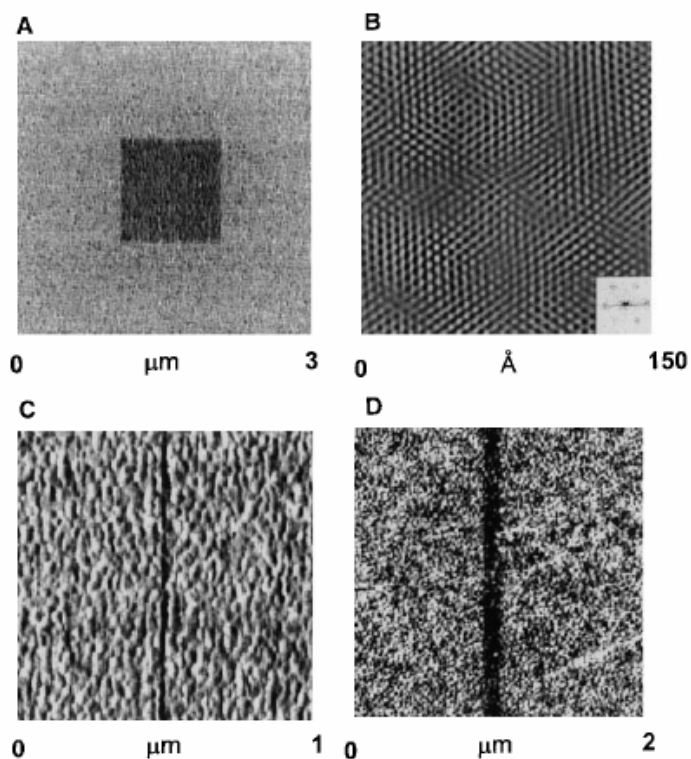


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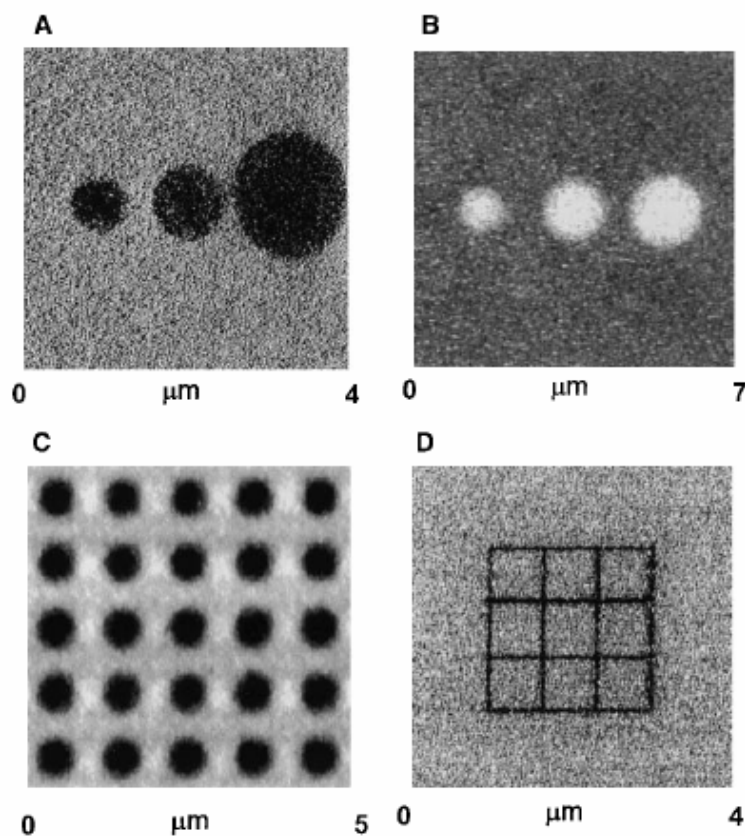
Dip-Pen Lithography

Fig. 2. (A) Lateral force image of a square of ODT measuring $1\ \mu\text{m}$ by $1\ \mu\text{m}$, deposited onto a Au substrate by DPN. This pattern was generated by scanning the $1\text{-}\mu\text{m}^2$ area at a scan rate of 1 Hz for a period of 10 min at a relative humidity of 39%. Then the scan size was increased to $3\ \mu\text{m}$, and the scan rate was increased to 4 Hz while the image was recorded. The faster scan rate prevents ODT transport. (B) Lattice-resolved, lateral force image of an ODT SAM deposited onto Au(111)/mica by DPN. The image has been filtered with a fast Fourier transform (FFT), and the FFT of the raw data is shown in the lower right insert. The monolayer was generated by scanning a $1000\ \text{\AA}$ square area of the Au(111)/mica five times at a rate of 9 Hz at 39% relative humidity. (C) Lateral force image of a 30-nm-wide line ($3\ \mu\text{m}$ long) deposited onto Au/mica by DPN. The line was generated by scanning the tip in a vertical line repeatedly for 5 min at a scan rate of 1 Hz. (D) Lateral force image of a 100-nm line deposited on Au by DPN. The method of depositing this line is analogous to that used to generate the image in (C), but the writing time was 1.5 min. In all images, darker regions correspond to areas of relatively lower friction.



Dip-Pen Lithography

Fig. 3. (A) Lateral force image of an Au substrate after an AFM tip, which was coated with ODT, had been in contact with the substrate for 2, 4, and 16 min (left to right); the relative humidity was held constant at 45%, and the image was recorded at a scan rate of 4 Hz. (B) Lateral force image of dots of 16-mercaptohexadecanoic acid on a Au substrate. To generate the dots, an AFM tip coated with 16-mercaptohexadecanoic acid was held on the Au substrate for 10, 20, and 40 s (left to right). The relative humidity was 35%. The images show that the transport properties of 16-mercaptohexadecanoic acid and of ODT differ substantially. (C) Lateral force image of an array of dots generated by DPN. Each dot was generated by holding an ODT-coated tip in contact with the surface for ~20 s. Writing and recording conditions were the same as in (A). (D) Lateral force image of a molecule-based grid. Each line is 100 nm in width and 2 μm in length and required 1.5 min to write.



Dip-pen Lithography

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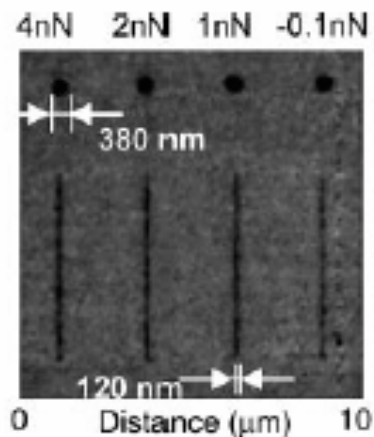
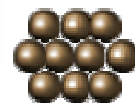
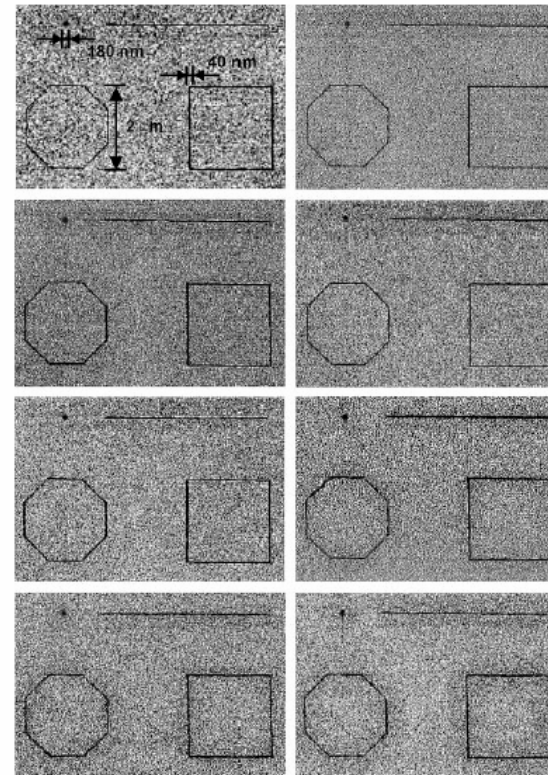


Fig. 1. Lateral force microscopy (LFM) images of ODT monolayer nanodot and line features on gold generated by the same tip but under different tip-substrate contact forces. Feature sizes vary less than 10%.

Fig. 4. LFM images of eight identical patterns generated with one imaging tip and eight writing tips coated with ODT molecules.



Protein Nanoarrays Generated By Dip-Pen Nanolithography

Ki-Bum Lee,¹ So-Jung Park,¹ Chad A. Mirkin,^{1*} Jennifer C. Smith,²
Milan Mrksich^{2*}

1 MARCH 2002 VOL 295 SCIENCE

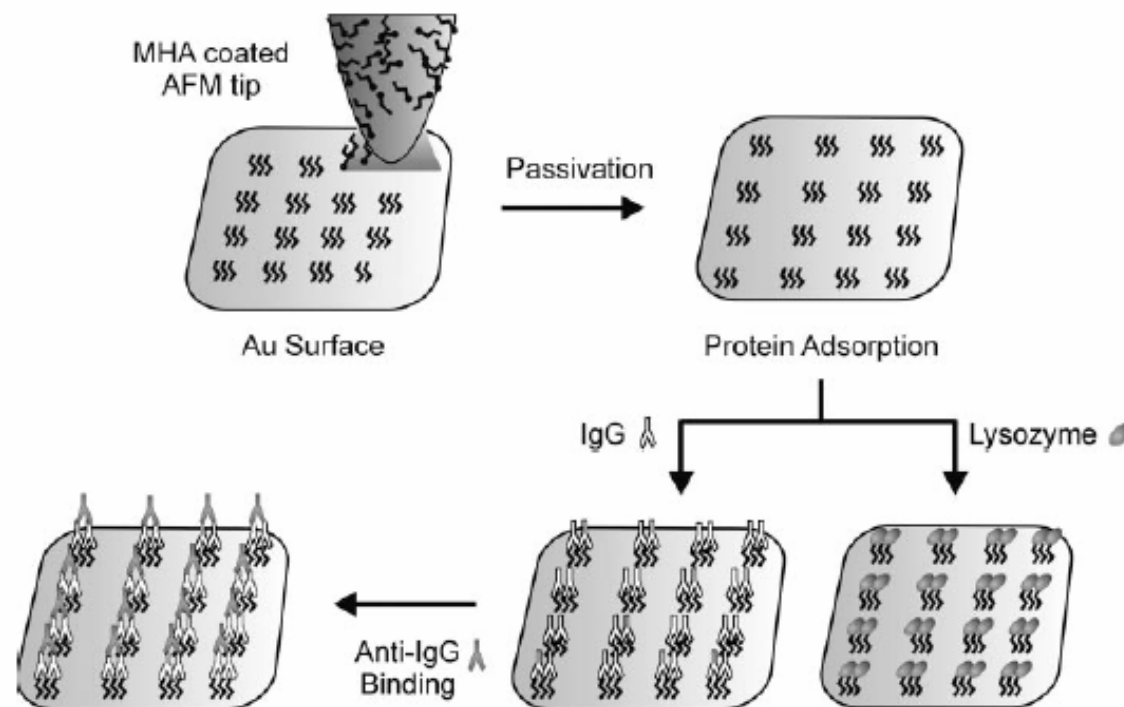
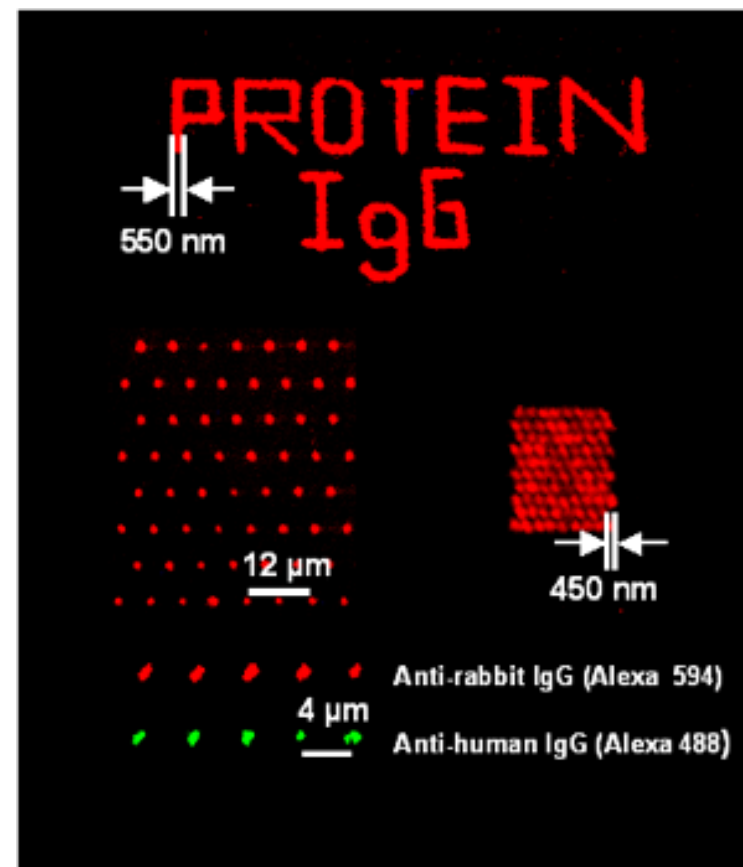


Fig. 2. Diagram of proof-of-concept experiments, in which proteins were absorbed on preformed MHA patterns. The resulting protein arrays were then characterized by AFM.



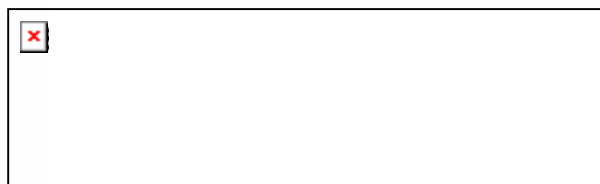
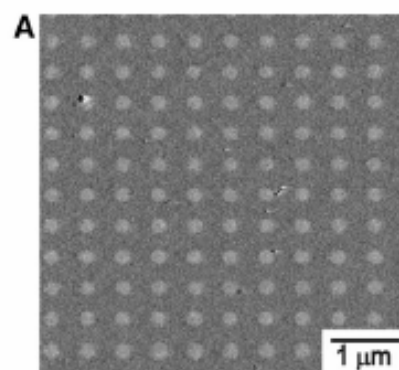
Protein Nanoarrays Generated By Dip-Pen Nanolithography

Ki-Bum Lee,¹ So-Jung Park,¹ Chad A. Mirkin,^{1*} Jennifer C. Smith,²
Milan Mrksich^{2*}

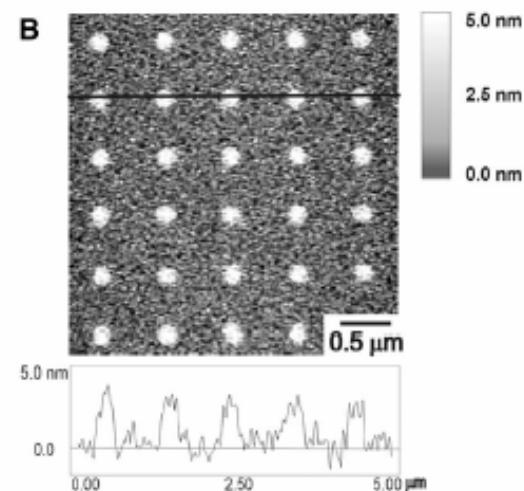
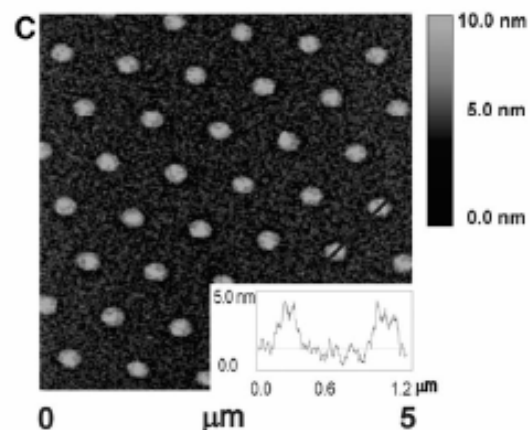


Protein Nanoarrays Generated By Dip-Pen Nanolithography

Ki-Bum Lee,¹ So-Jung Park,¹ Chad A. Mirkin,^{1*} Jennifer C. Smith,²
Milan Mrksich^{2*}



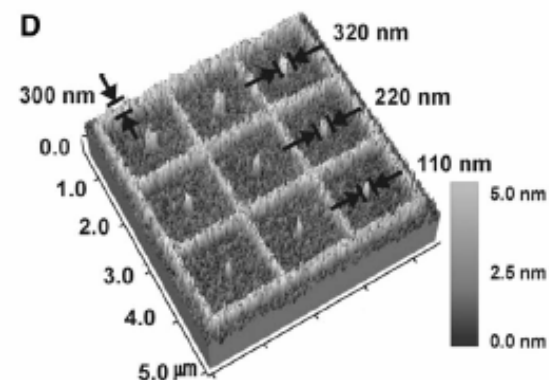
5.



n

n

n



Protein Nanoarrays Generated By Dip-Pen Nanolithography

Ki-Bum Lee,¹ So-Jung Park,¹ Chad A. Mirkin,^{1*} Jennifer C. Smith,²
Milan Mrksich^{2*}

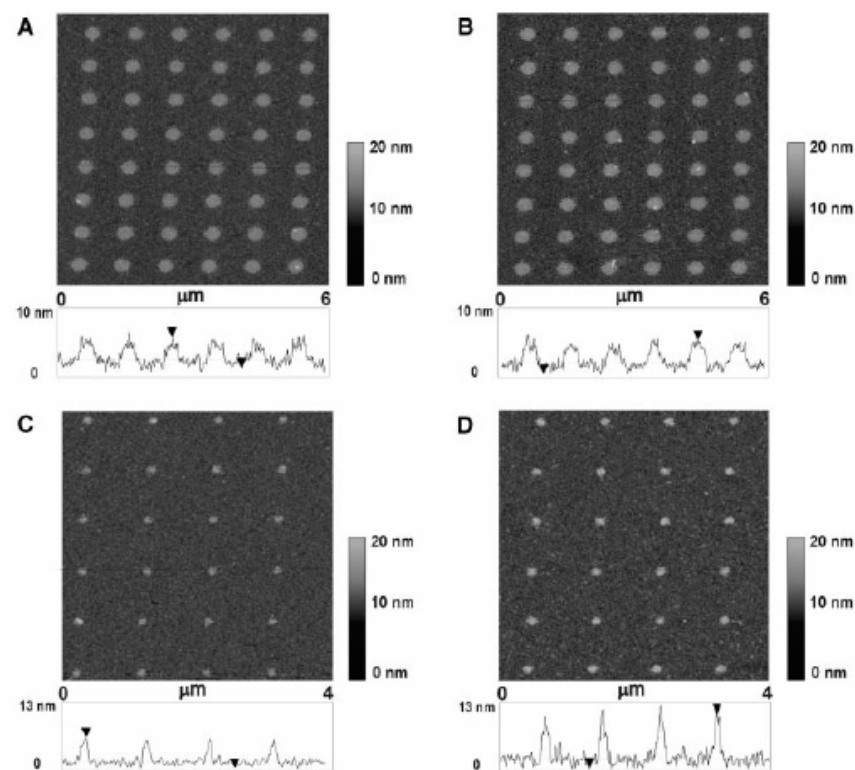


Fig. 3. AFM tapping mode image and height profile of rabbit IgG assembled onto an MHA dot array generated by DPN before (A) and after (B) exposure to a solution containing lysozyme, Retronectin, goat/sheep anti-IgG, and human anti-IgG. An IgG nanoarray before (C) and after (D) treatment with a solution containing lysozyme, goat/sheep anti-IgG, human anti-IgG, and rabbit anti-IgG. All images were taken at a 0.5-Hz scan rate in tapping mode.



Protein Nanoarrays Generated By Dip-Pen Nanolithography

Ki-Bum Lee,¹ So-Jung Park,¹ Chad A. Mirkin,^{1*} Jennifer C. Smith,²
Milan Mrksich^{2*}

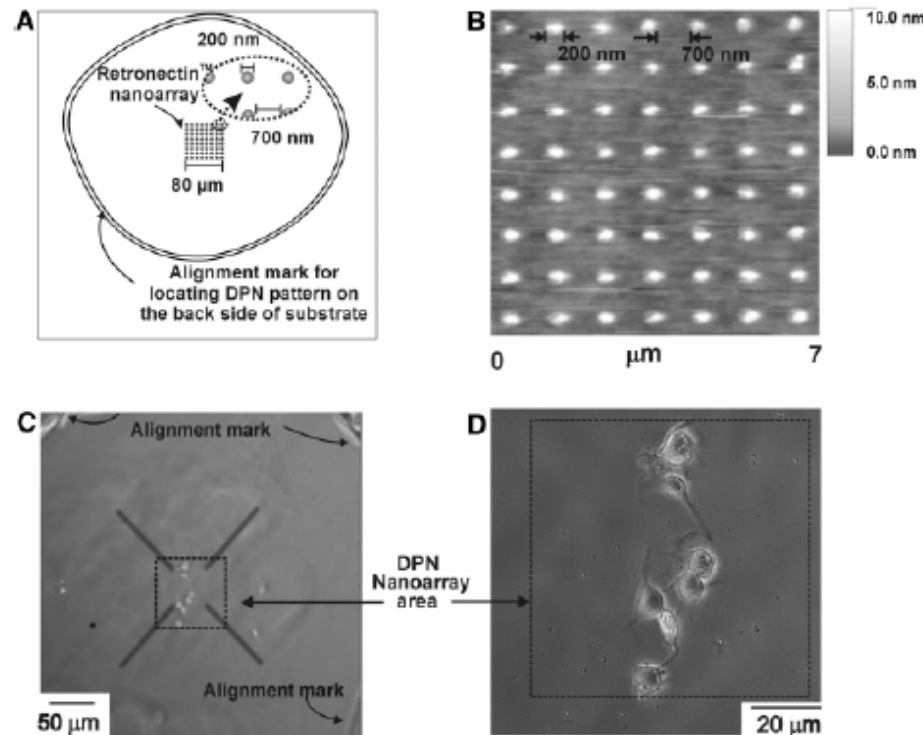


Fig. 4. (A) Diagram describing the cell adhesion experiment on the DPN-generated pattern. The total patterned area is 6400 μm². The alignment marks were generated by scratching a circle into the backside of the Au-coated glass substrate. (B) Topography image (contact mode) of the Retronectin protein array. Imaging conditions were the same as in Fig. 1B. (C) Large-scale optical microscope image showing the localization of cells in the nanopatterned area. (D) Higher resolution optical image of the nanopatterned area, showing intact cells.



Dip-Pen Array

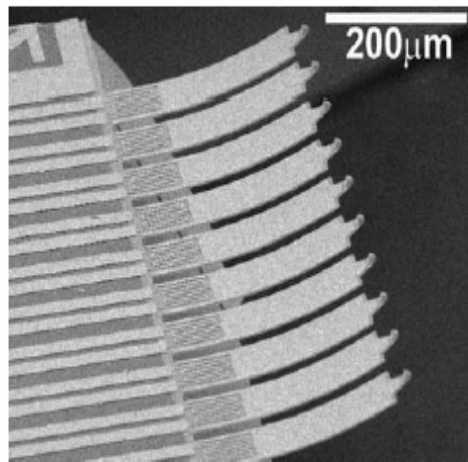


FIG. 2. An array of ten thermally actuated DPN probes showing the power lead and heater layout. Each probe is 300 μm long, 80 μm wide, and 1.3 μm thick.

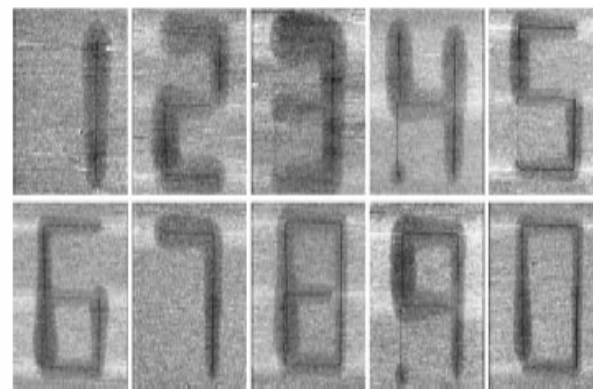
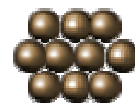
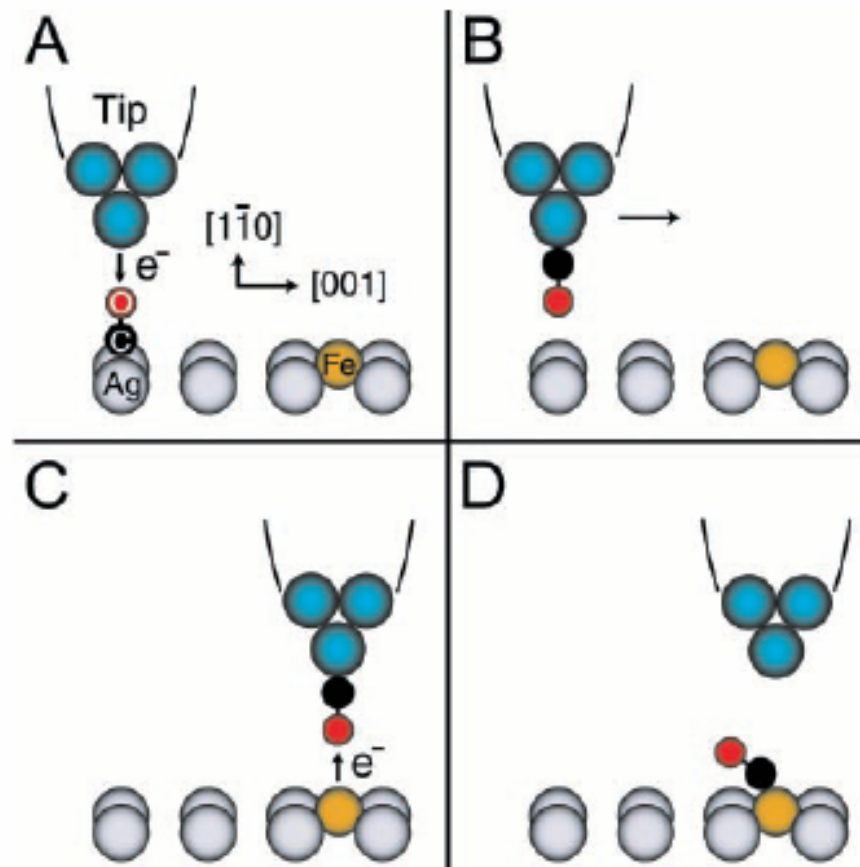


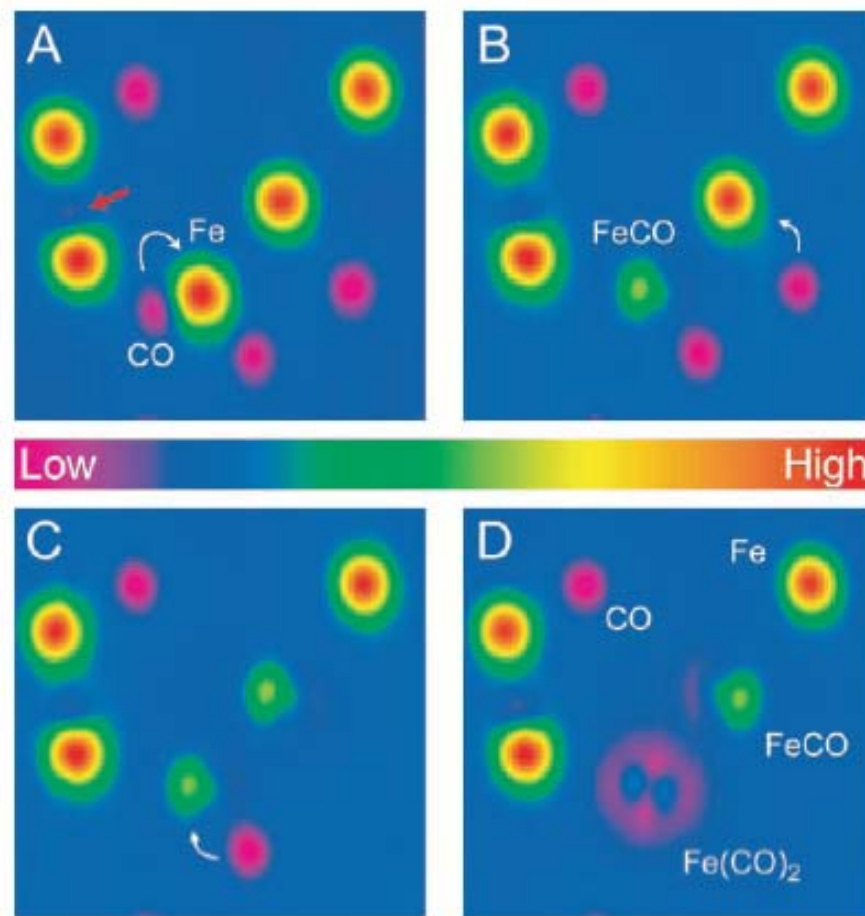
FIG. 4. LFM scans, 8 μm square, of ten simultaneously generated ODT patterns on a gold surface. Each numeral is 6 μm tall, 4 μm wide, and was written at 1 $\mu\text{m/s}$.



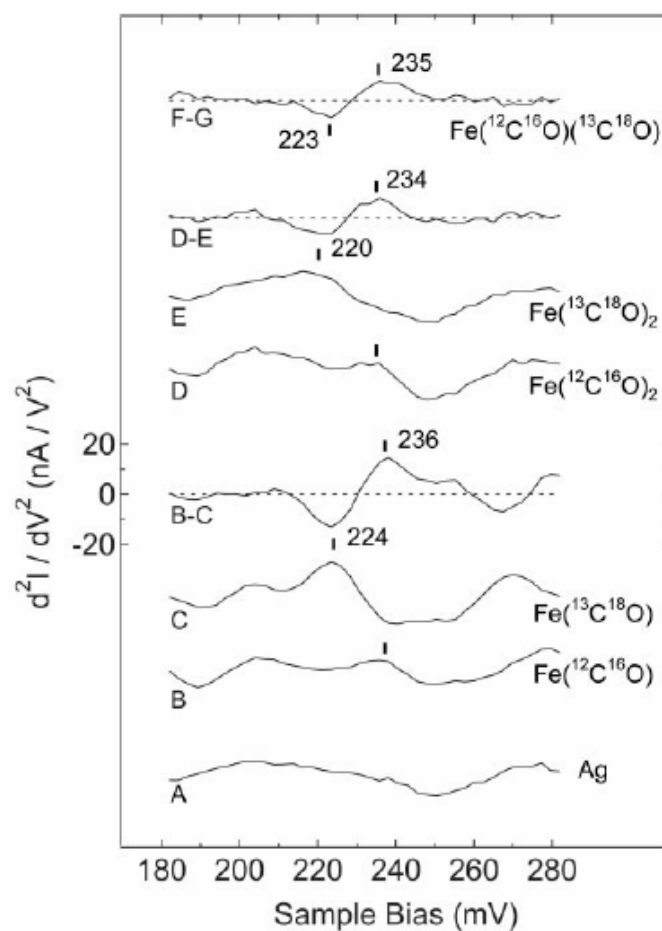
Ultimate STM Lithography



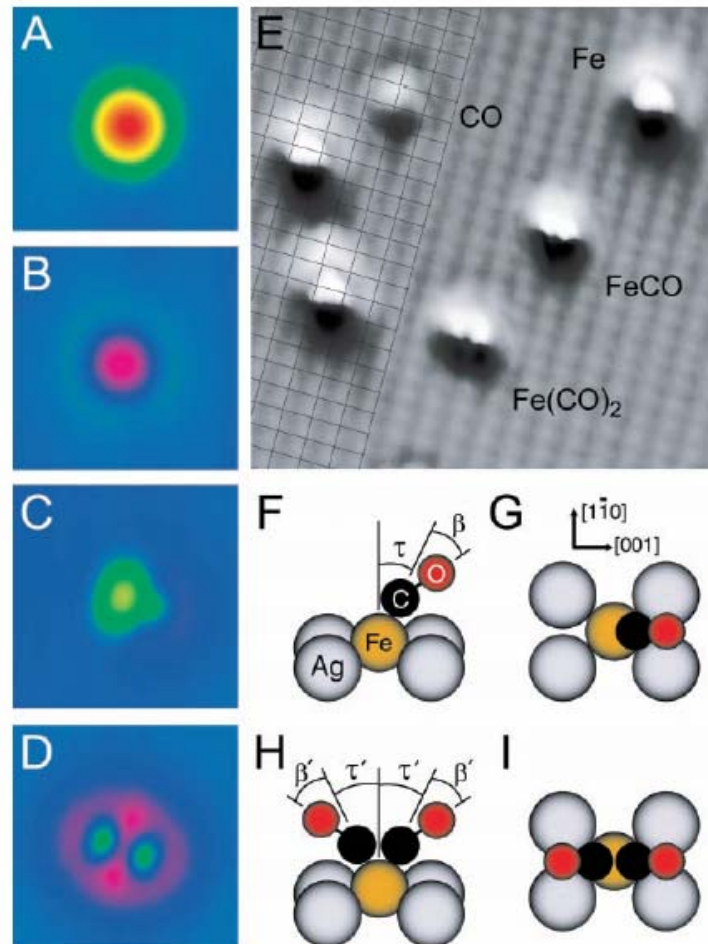
Single Atomic Manipulation



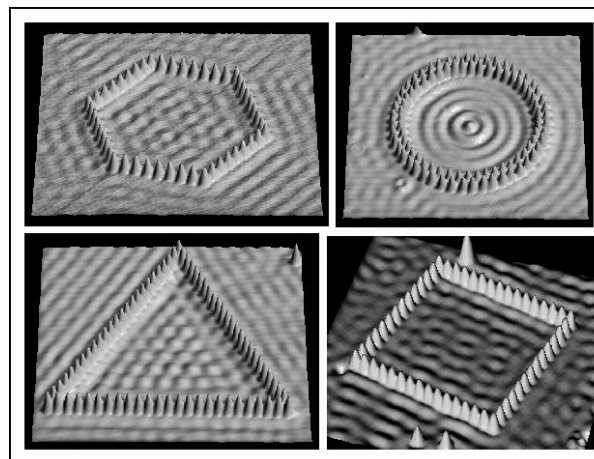
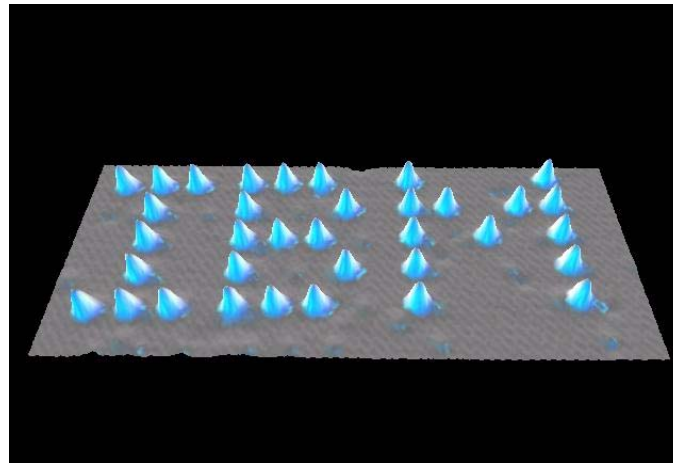
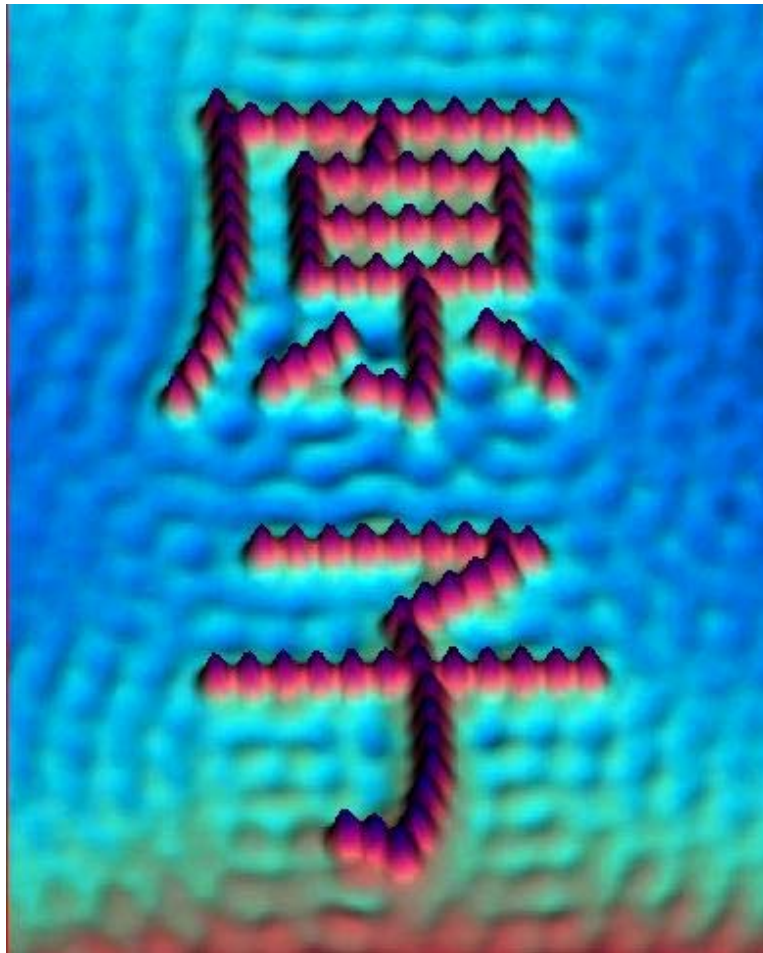
Single Molecular Vibrational Spectra by STM



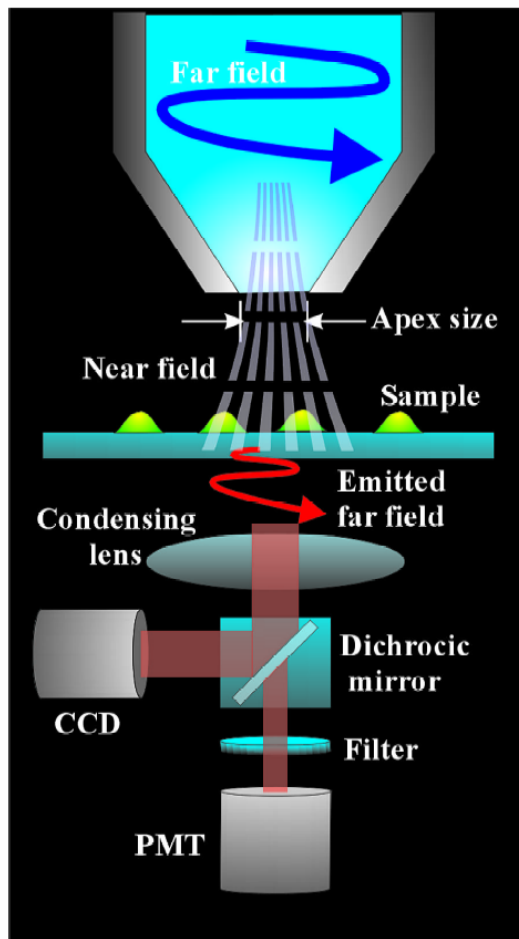
Building Molecule Step by Step



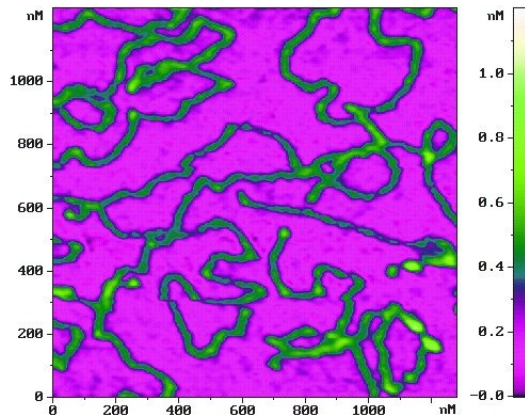
Atomic Manipulation



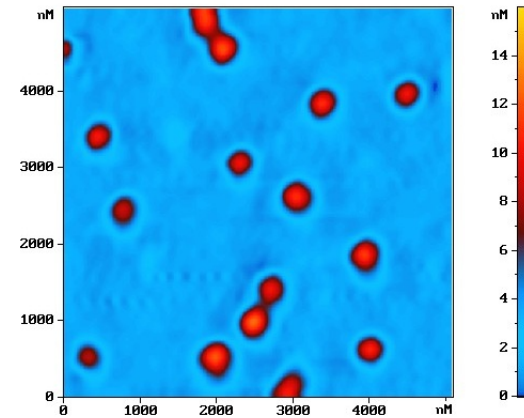
Near-Field Microscope



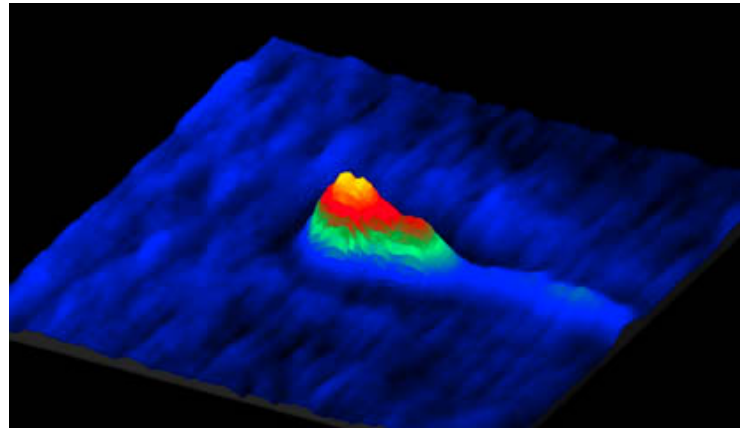
Near-Field Images



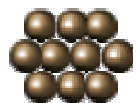
DNA



Nanosphere



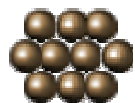
Sperm



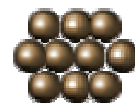
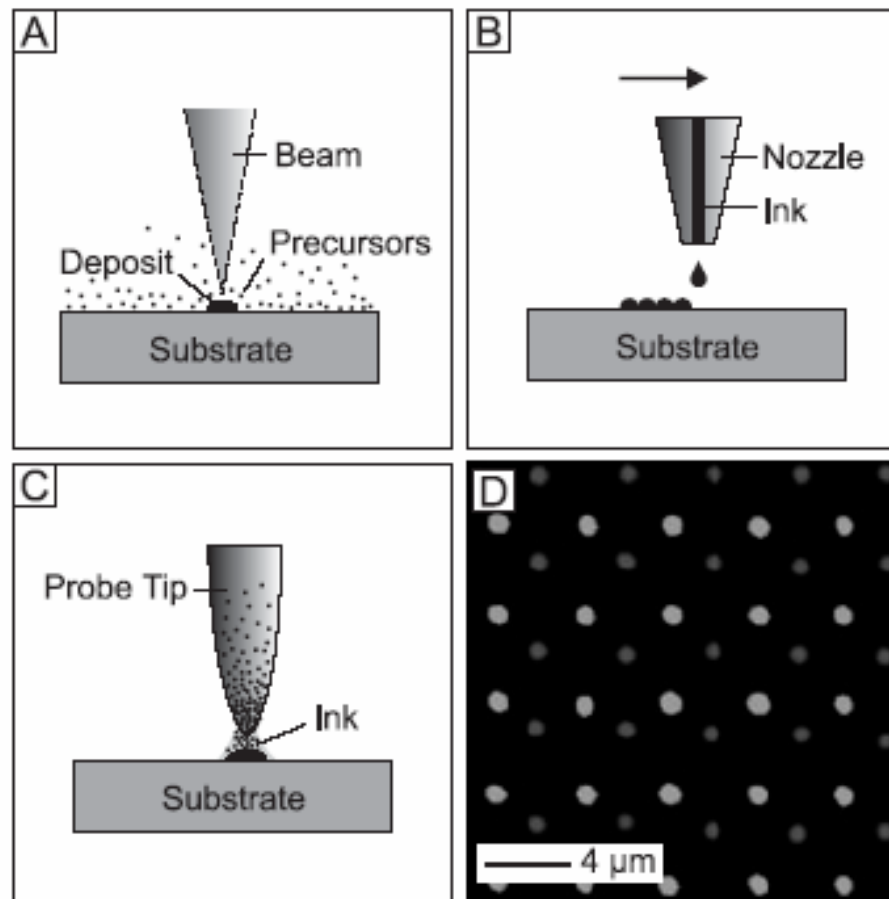
Near-Field Lithography



Figure 2. TappingMode image of a lithography test pattern. The Aurora-3 used nanolithography software to write into S1805 photoresist. 25 μ m scan.



Add on Writing



Direct Writing 3D

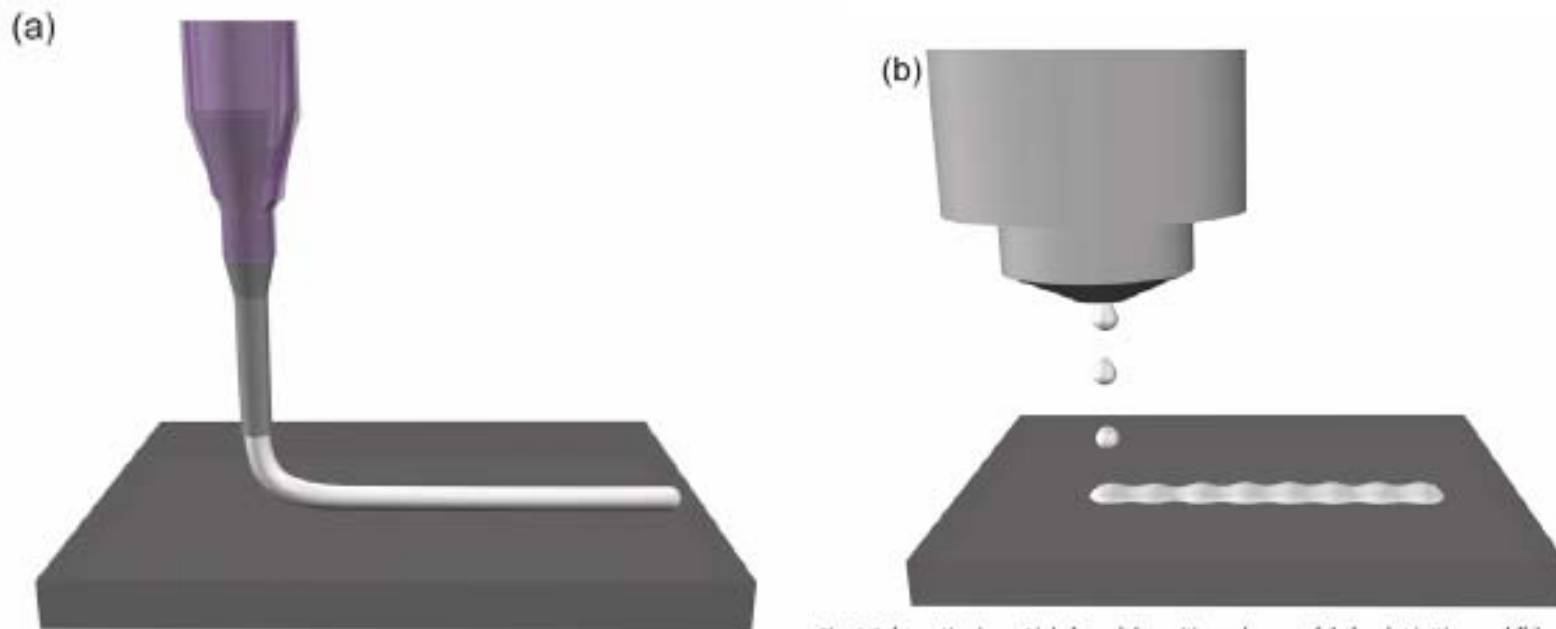
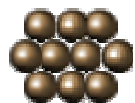


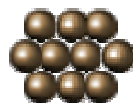
Fig. 1 Schematic view of ink-based deposition schemes: (a) droplet jetting and (b) continuous filament writing. (Reprinted from ³¹, ©2002 with permission from Elsevier Ltd.)

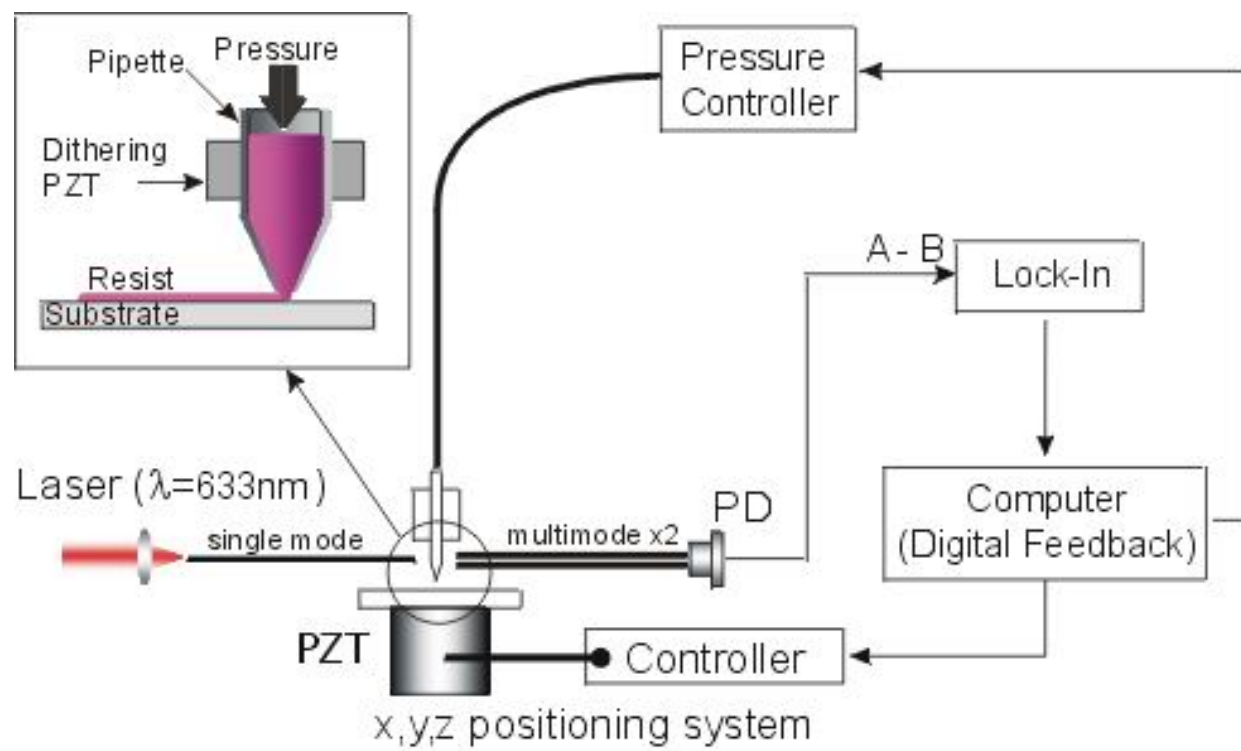


Direct Writing

Table 1 Capabilities of ink-writing techniques.

Technique	Ink design	Minimum printed feature size	3-D periodic structures
Robotic Deposition ¹¹	Concentrated colloidal gel ^{13,14}	200 μm diameter	Yes
•	Concentrated nanoparticle gel ¹⁵	100 μm diameter	Yes
•	Viscous polymer solution ¹⁶⁻¹⁸	200 μm diameter	Yes
•	Concentrated polyelectrolyte complexes ³	<1 μm	Yes
Three-dimensional printing ¹⁹	Binder solution printed on powder bed	170 μm lateral, 45 μm depth	Yes
Ink-jet printing ²⁰	Dilute fluid ²¹	20 μm lateral, 100 nm height	No
•	Concentrated fluid (max. solids ~ 40%) ²²⁻²⁵	70 μm lateral, < 1 μm height	No
Fused deposition	Thermoplastic polymer melt ²⁵	100 μm diameter	Yes
•	Particle-filled polymer melt ²⁷ (max. solids ~ 50%)	100 μm diameter	Yes
Micropen writing ²⁶	Concentrated, shear-thinning colloidal fluid	25 μm diameter	No
Dip-pen nanolithography ²⁹	Dilute fluid	20 nm	No
Scanning probe contact printing ³⁰	Dilute fluid	<500 nm	No





Direct Writing

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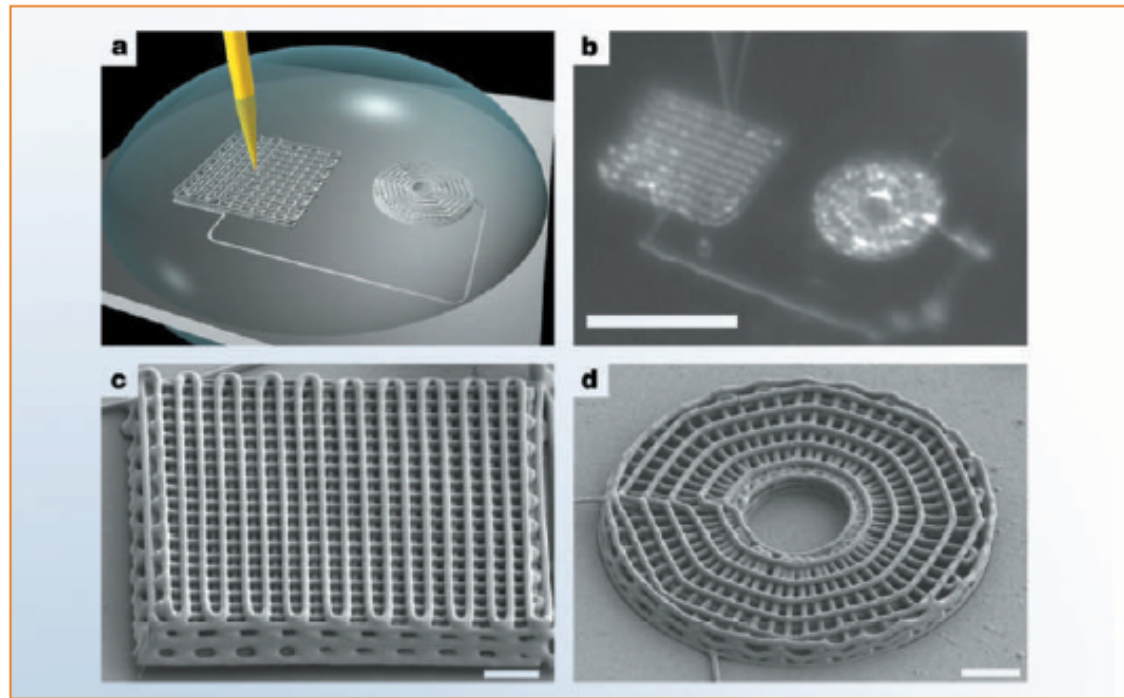
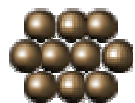


Figure 1 Direct-write assembly of three-dimensional microperiodic structures. **a**, The ink-deposition process (not drawn to scale). A concentrated polyelectrolyte ink is housed in a syringe (yellow) immersed in a coagulation reservoir (grey hemispherical drop) and deposited on to a glass substrate (light grey). **b**, Optical image acquired *in situ* during deposition reveals the features drawn in **a**, including the deposition nozzle that is patterning a three-dimensional lattice, as well as a completed radial array. This image is blurred by the reservoir (scale bar: 100 μm). **c**, Three-dimensional periodic structure with a face-centred tetragonal geometry (filament diameter: 1 μm ; 10 layers; scale bar: 10 μm). **d**, Three-dimensional radial array (filament diameter: 1 μm ; 5 layers; scale bar: 10 μm).



Inkjet Printer



Figure 1. Microdrop Autodrop Platform (courtesy Microdrop GmbH, Germany).



Inkjet Printing

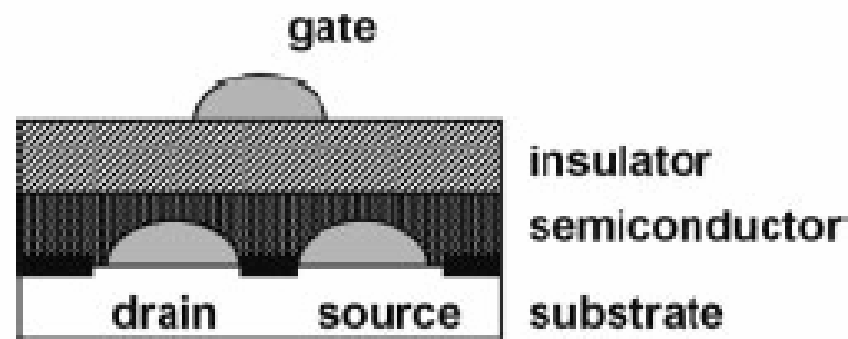
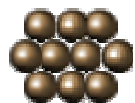


Figure 5. Schematic picture of inkjet printed all-polymer thin film transistor, as constructed by Sirringhaus et al. [41]. Source and drain electrode, consisting of PEDOT/PPS are inkjet printed on a pre-patterned surface. Two spin-coated layers of semiconducting and insulating polymer respectively cover the electrodes. Finally the gate electrode is printed on top.



Inkjet Printing

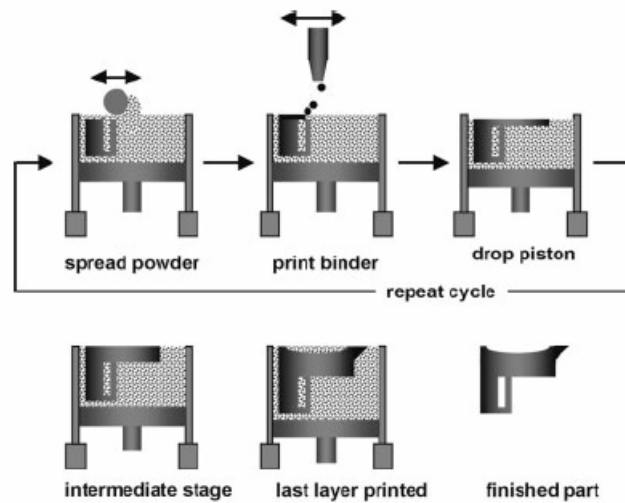


Figure 7. Schematic picture of the three-dimensional printing process.

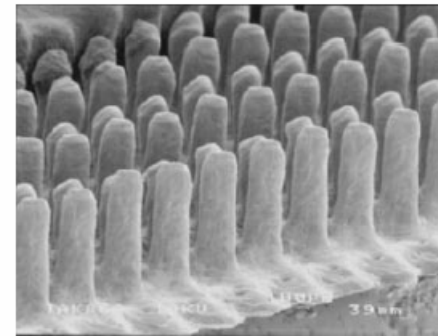


Figure 8. Electron microscopy picture of an array of pillars from lead zirconium titanate, inkjet printed by Bhatti et al. The pillars have an approximate height of 400 μm. (Reprinted from [59] with permission. Copyright 2001 Kluwer Academic Publishers).



Inkjet Printing

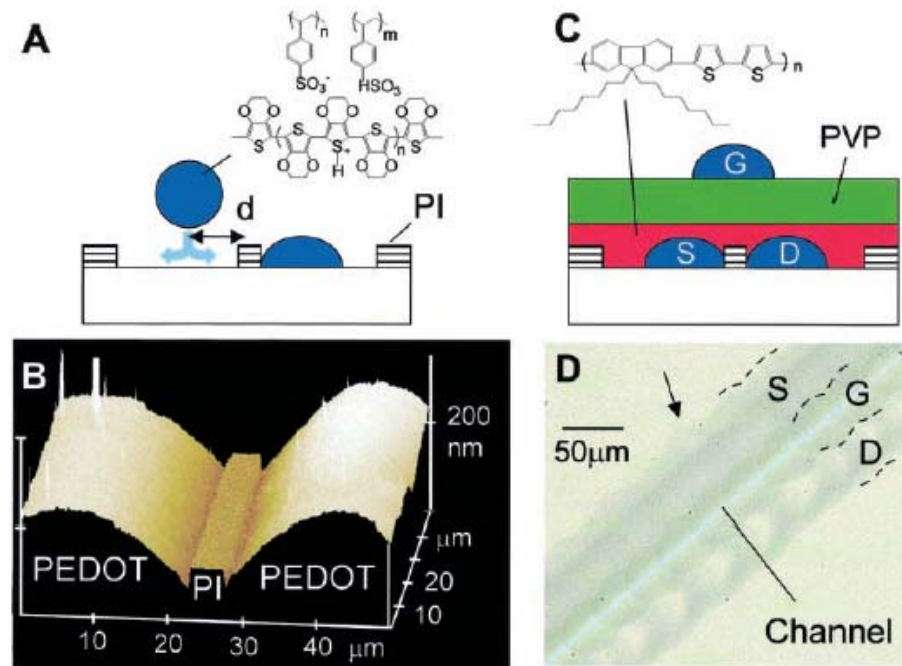
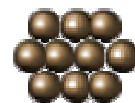
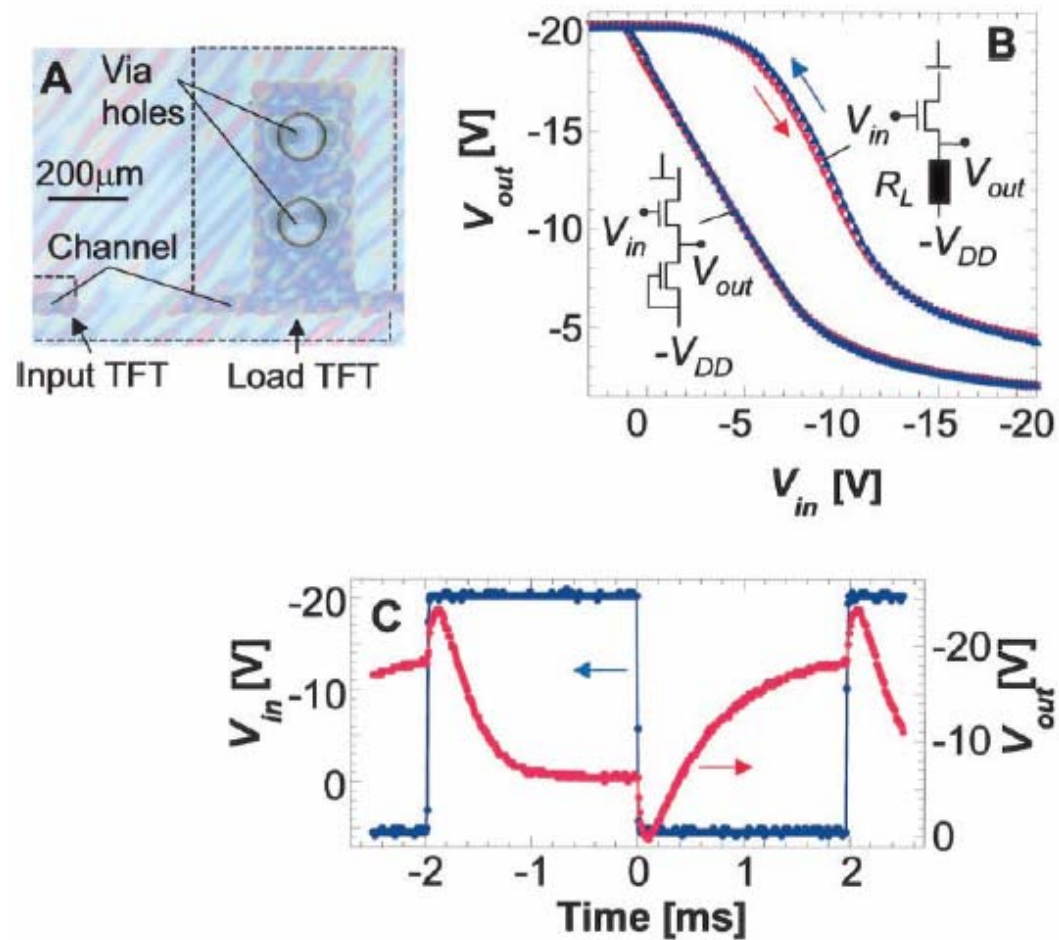


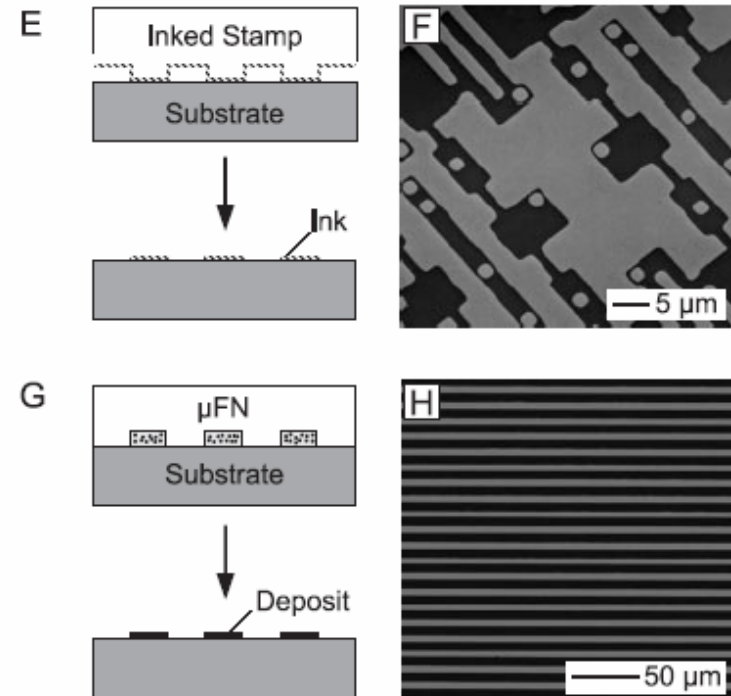
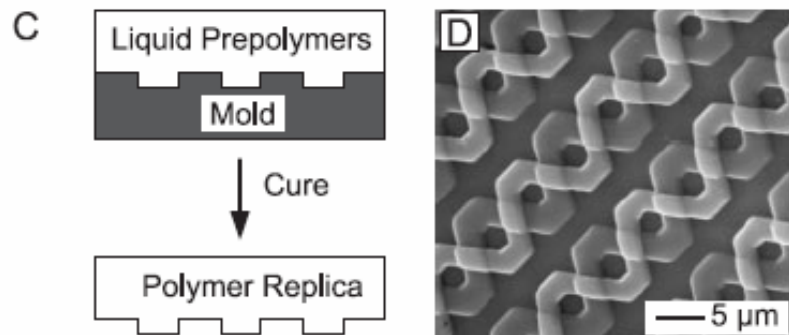
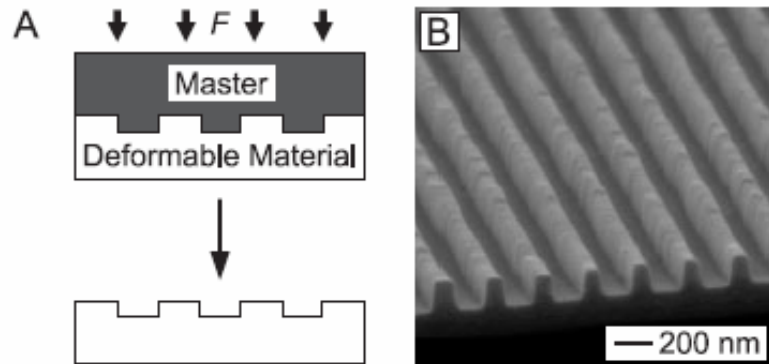
Fig. 1. (A) Schematic diagram of high-resolution IJP onto a prepatterned substrate. (B) AFM showing accurate alignment of inkjet-printed PEDOT/PSS source and drain electrodes separated by a repelling polyimide (PI) line with $L = 5 \mu\text{m}$. (C) Schematic diagram of the top-gate IJP TFT configuration with an F8T2 semiconducting layer (S, source; D, drain; and G, gate). (D) Optical micrograph of an IJP TFT ($L = 5 \mu\text{m}$). The image was taken under crossed polarizers so that the TFT channel appears bright blue because of the uniaxial monodomain alignment of the F8T2 polymer on top of rubbed polyimide. Unpolarized background illumination is used to make the contrast in the remaining areas visible, where the F8T2 film is in an isotropic multidomain configuration. The arrow indicates pronounced roughness of the unconfined PEDOT boundary.



Inkjet Printing

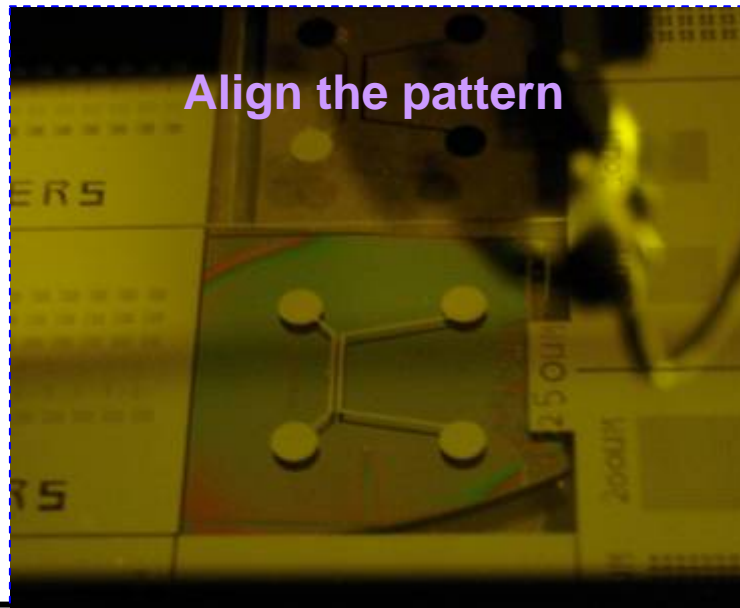
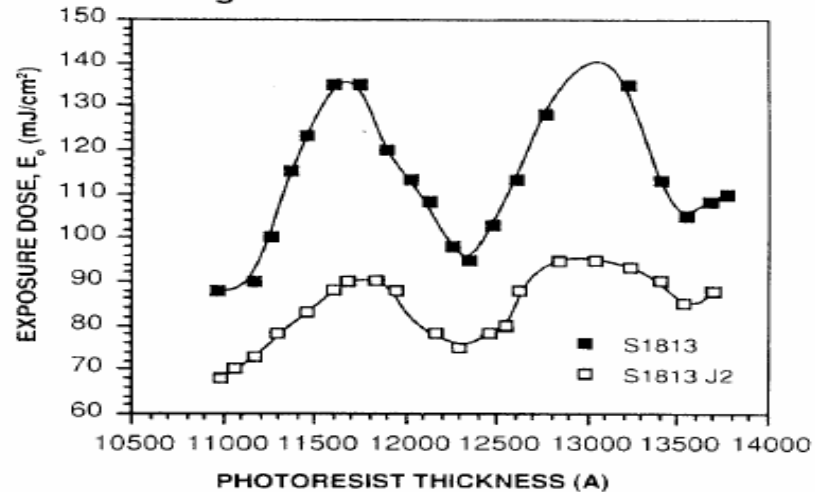


Replication

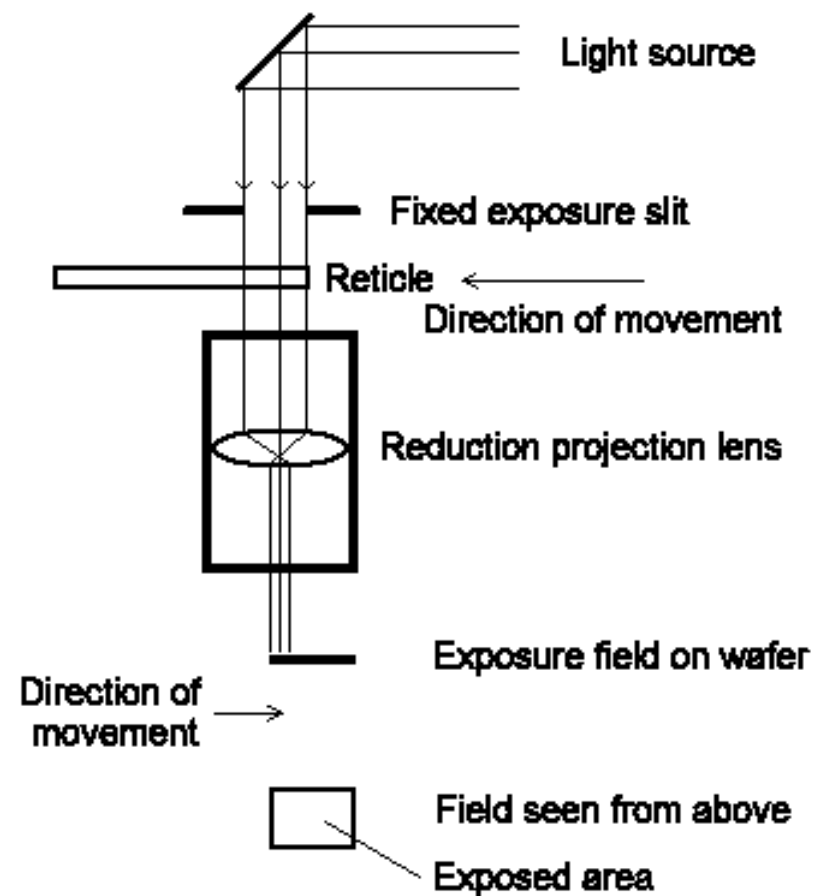
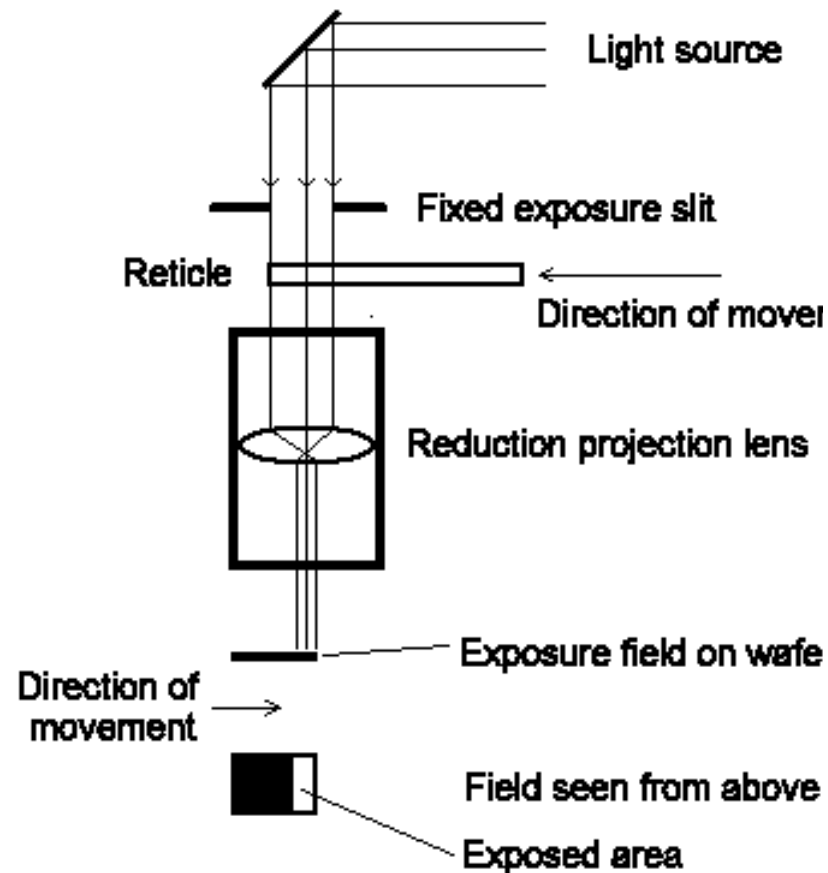


Align the pattern and Exposure

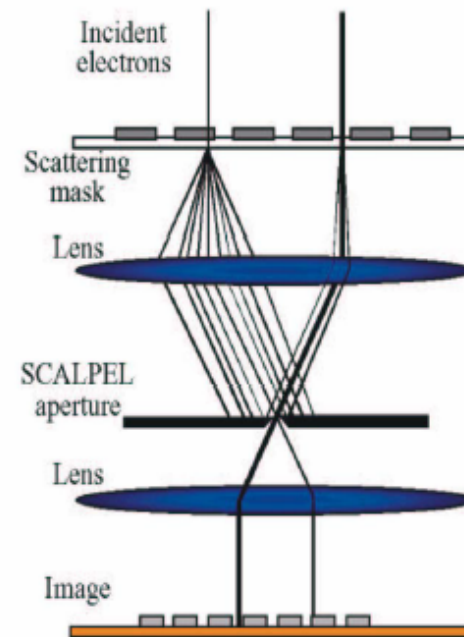
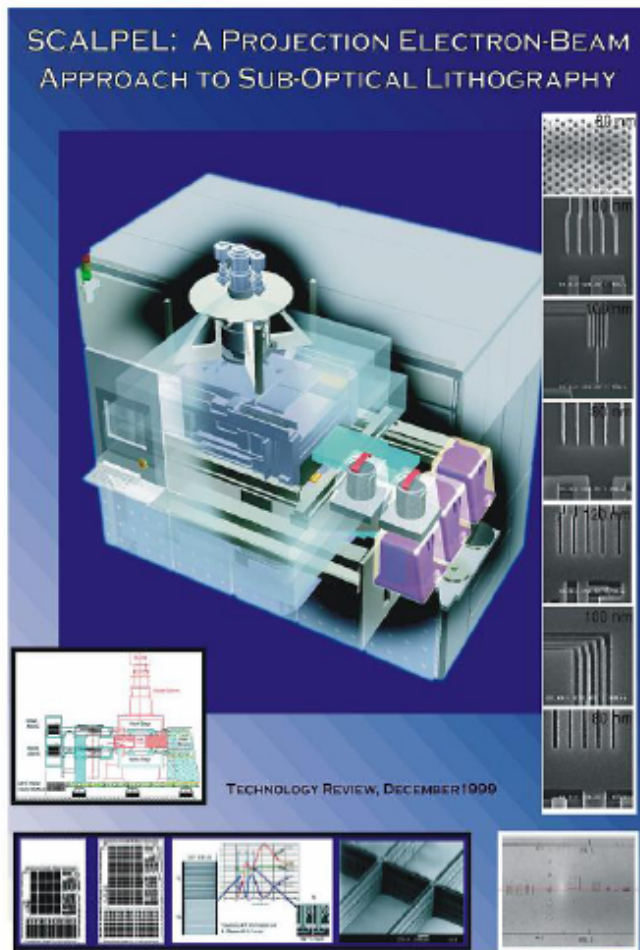
MICROPOSIT S1813 and S1813 J2 PHOTO RESISTS
Figure 4. Interference Curves



Stepper

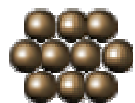


E-beam Projection



Bell Lab (1999)

There 'was' a consortium including Applied Materials, Inc. and ASM Lithography Holding N.V.; Lucent Technologies Inc.; Motorola, Semiconductor Products Sector; Samsung Electronics Co., Ltd.; and Texas Instruments Incorporated (TI).



Imprint Lithography with 25-Nanometer Resolution

Stephen Y. Chou; Peter R. Krauss; Preston J. Renstrom

Science, New Series, Volume 272, Issue 5258 (Apr. 5, 1996), 85-87.

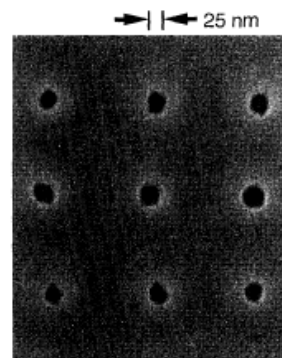


Fig. 2. SEM micrograph of a top view of holes 25 nm in diameter with a period of 120 nm, formed by compression molding into a PMMA film.

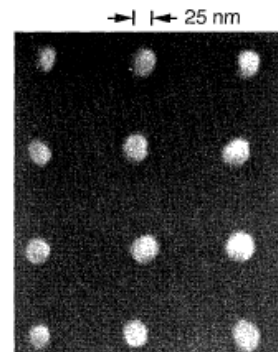


Fig. 5. SEM micrograph of the substrate in Fig. 2, after deposition of metal and a lift-off process. The diameter of the metal dots is 25 nm, the same as that of the original holes created in the PMMA.

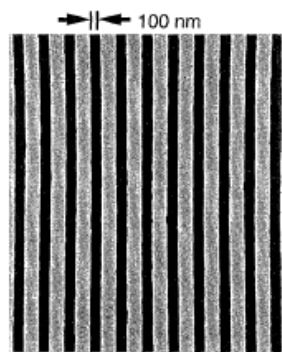


Fig. 3. SEM micrograph of a top view of trenches 100 nm wide with a period of 250 nm, formed by compression molding into a PMMA film.

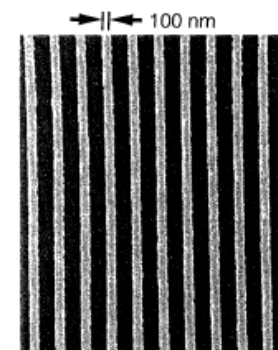
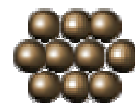
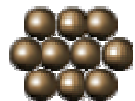
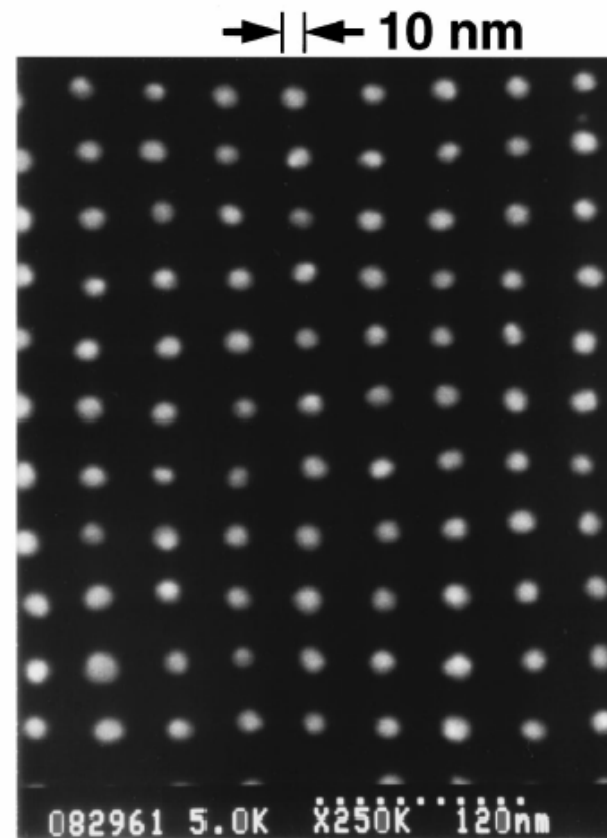
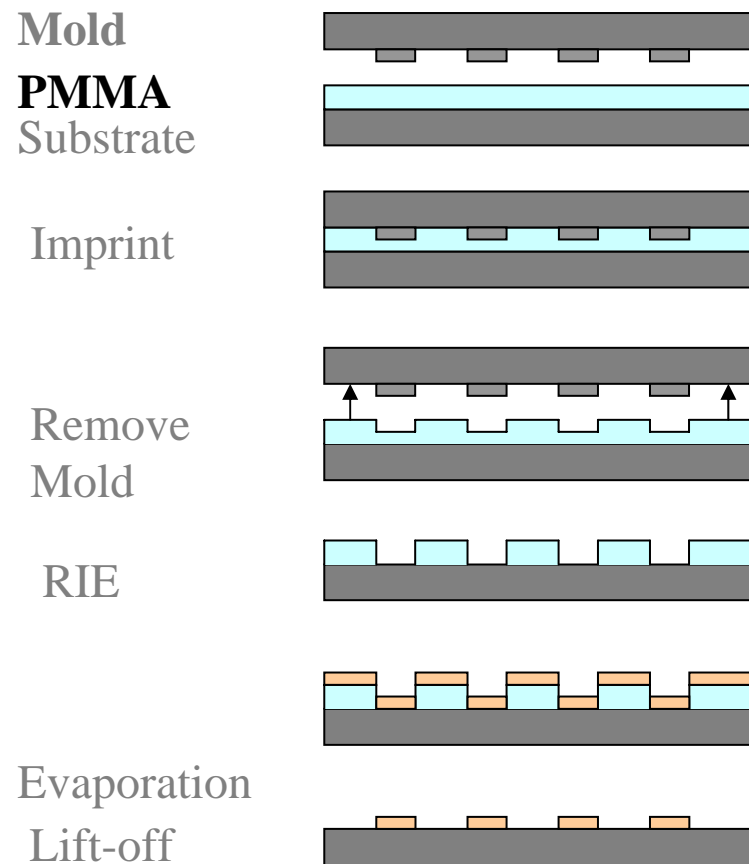


Fig. 6. SEM micrograph of the substrate in Fig. 3, after deposition of metal and a lift-off process. The metal linewidth is 100 nm, the same as the width of the original PMMA trenches.



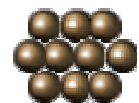
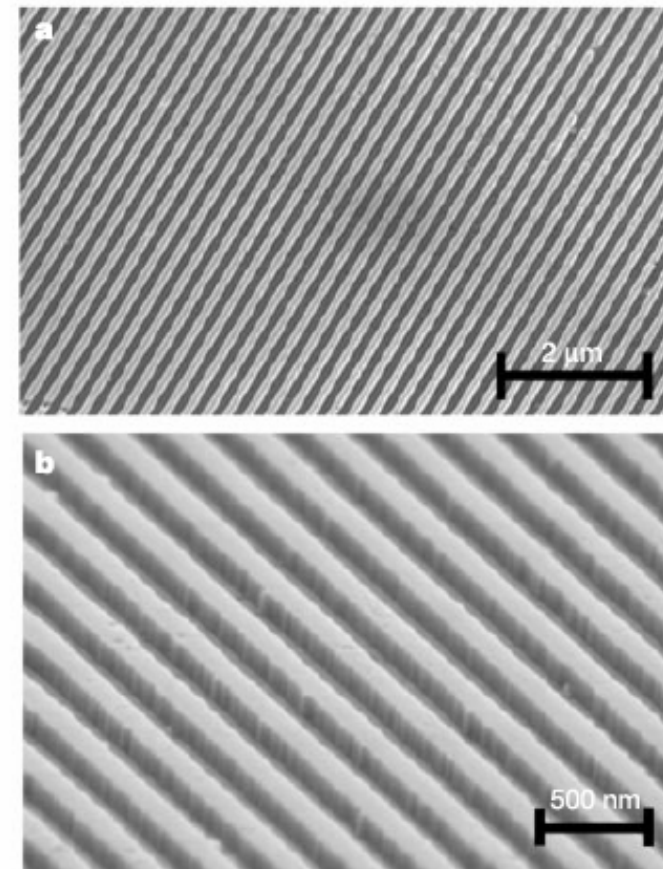
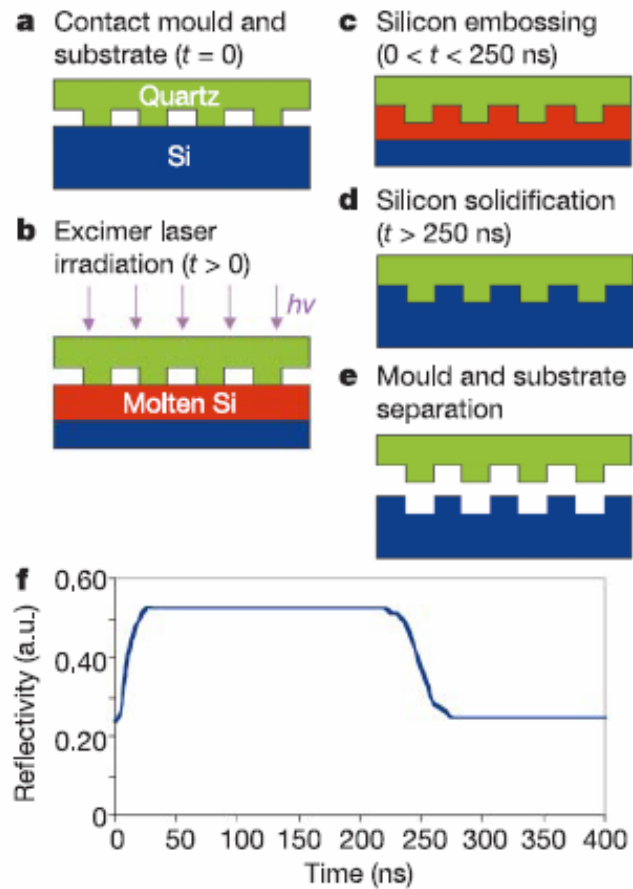
Nanoimprint Lithography



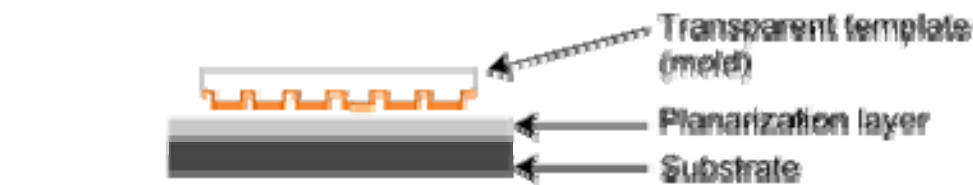
Ultrafast and direct imprint of nanostructures in silicon

NATURE | VOL 417 | 20 JUNE 2002 |

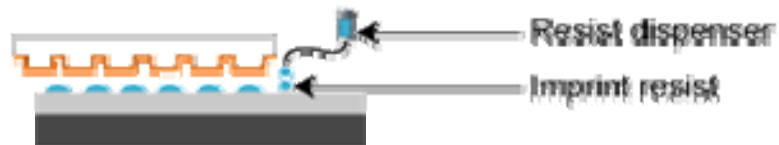
Stephen Y. Chou*, Chris Keimel & Jian Gu



Step and Flash Imprint Lithography



Step 1: Orient template and substrate



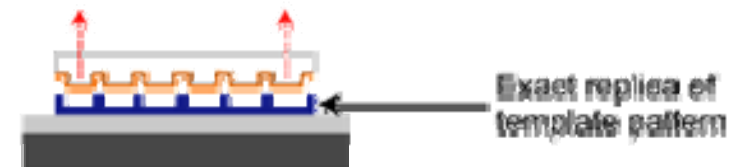
Step 2: Dispense drops of liquid imprint resist



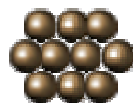
Step 3: Lower template and fill pattern



Step 4: Polymerize imprint fluid with UV exposure



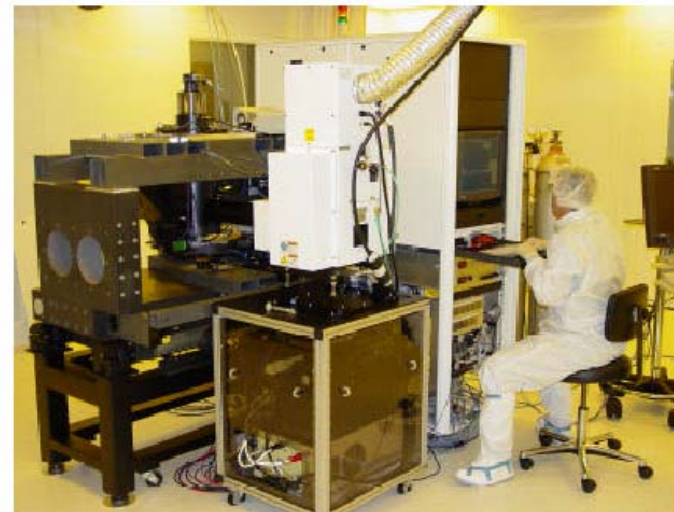
Step 5: Separate template from substrate



Nanoimprintors



NX-2000, Nanoimprintor, Nanonex



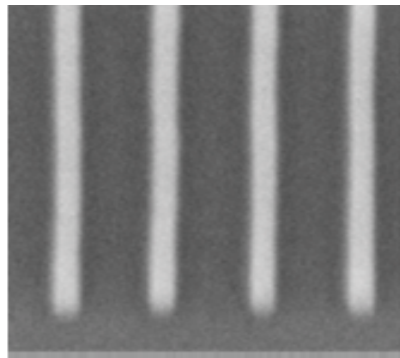

IMPRIO
100



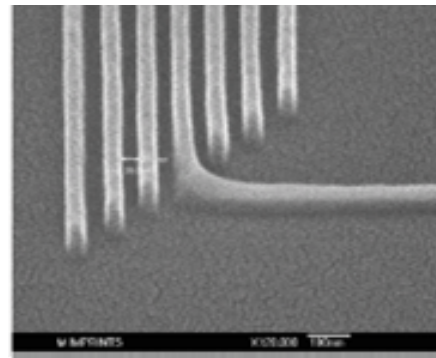
- Resolution: Sub-50 nanometers, imprint template (mold) limited.
- Alignment: < 500 nm, 3σ (X, Y, and Rotation).
- Flexibility: Handles up to 8 inch wafers, including fragile substrates.
- Field size: 25 x 25 mm full active print area, 100 μ m street width.



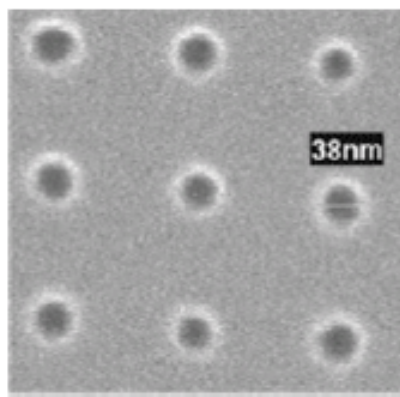
Imprinting Result



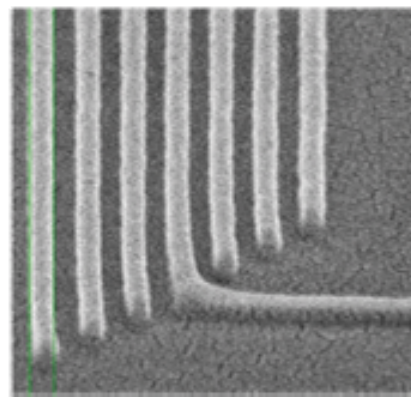
Imprinted 20 nm isolated lines



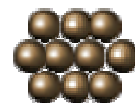
Imprinted 30 nm dense lines



Imprinted sub-40 nm contacts



Imprinted 50 nm dense lines



Challenges

- Mask Fabrication (1:1)
- Lift-off process
- Resist
- Mask Design

