

Consequences of Implanting and Surface Mixing During Si and SiO₂ Plasma Etching*

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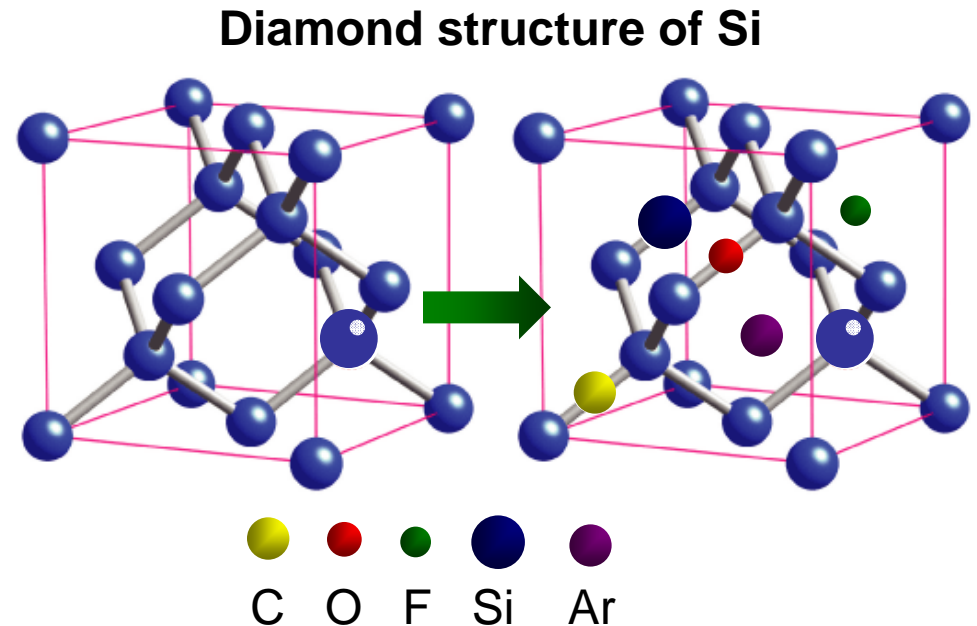
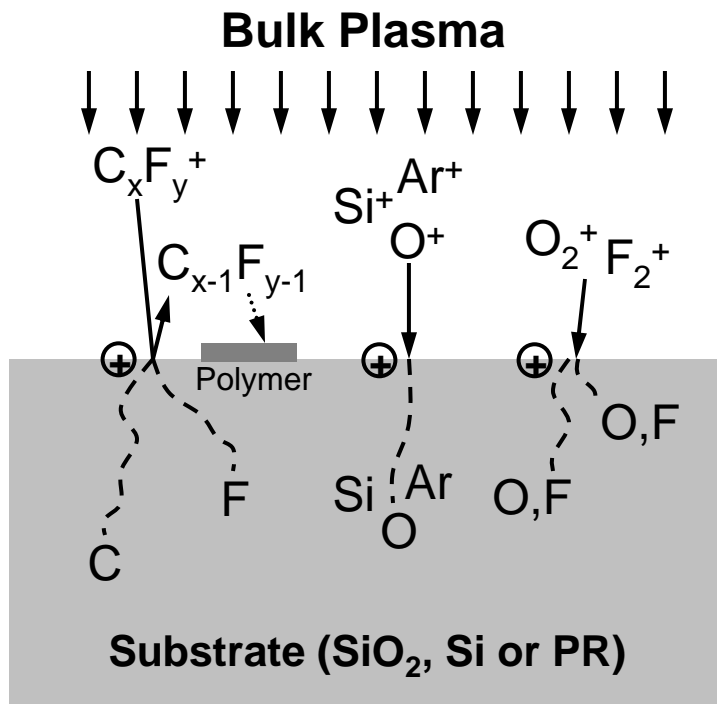
36th ICOPS, June 2009, San Diego, CA

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AGENDA

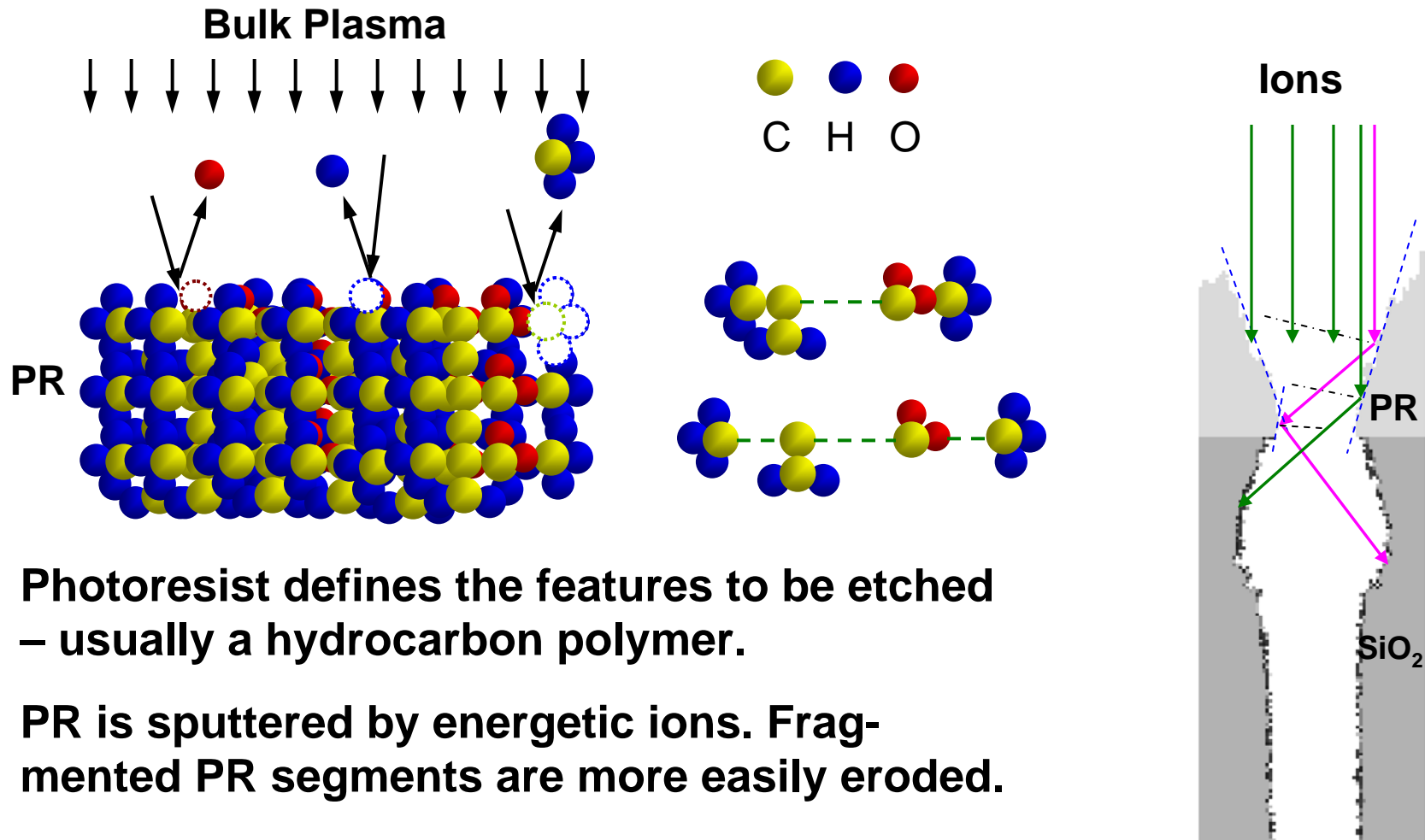
- **Implanting, mixing and photoresist (PR) erosion**
- **Processes**
 - **Molecular ion dissociation on surfaces**
 - **Small ion penetration and mixing**
 - **PR cross-linking**
- **Description of Model**
- **Scaling of Mixing and Implantation**
- **Concluding Remarks**

IMPLANTATION AND MIXING DURING PLASMA ETCHING



- Small ions accelerated by the sheath implant into the wafer surface forming weakly bonded or interstitially trapped species.
- Implanting causes surface mixing which produces damage during plasma etching.

PHOTORESIST (PR) EROSION

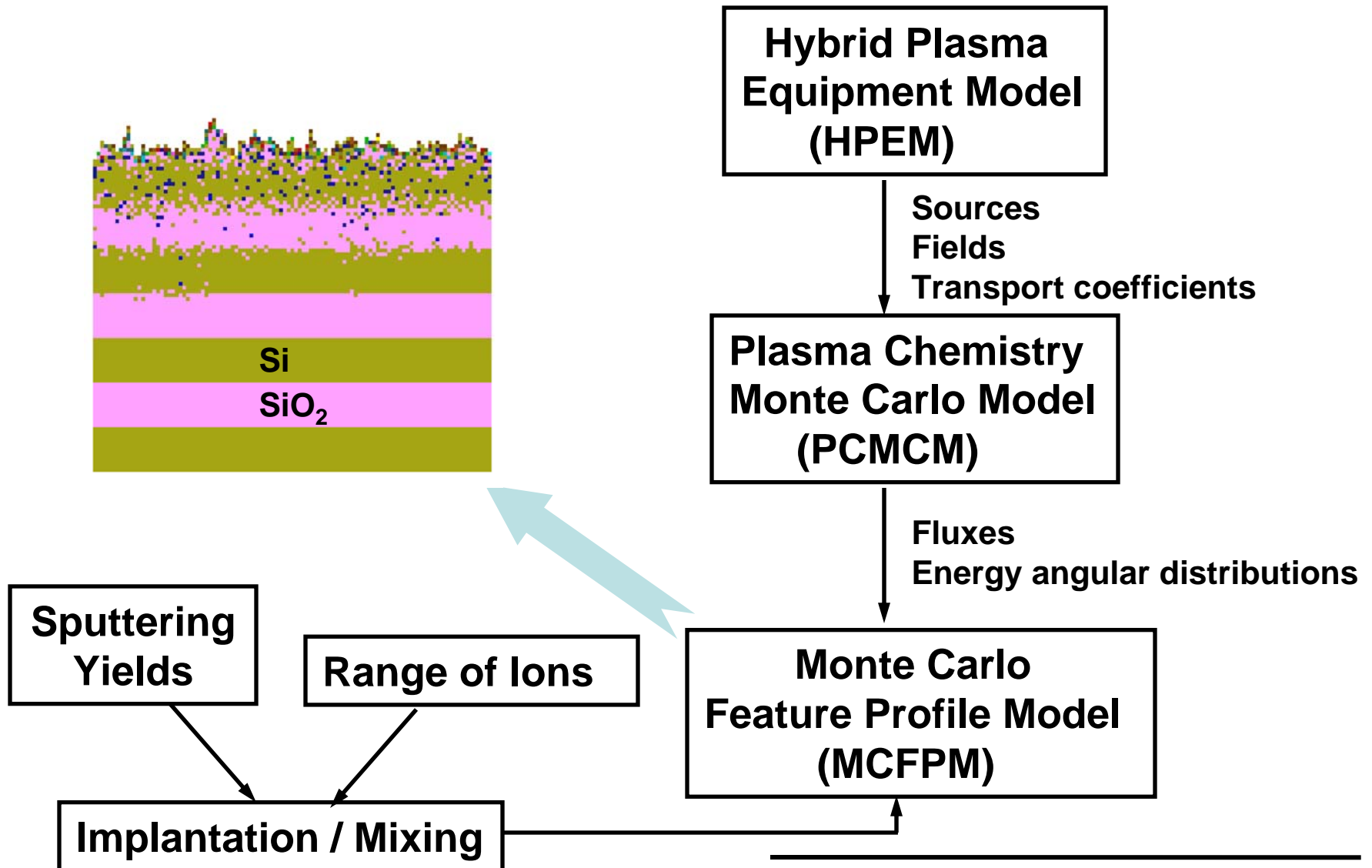


- Photoresist defines the features to be etched – usually a hydrocarbon polymer.
- PR is sputtered by energetic ions. Fragmented PR segments are more easily eroded.
- Profile of high aspect ratio (HAR) features can be modified due to PR erosion.

APPROACHES AND GOALS OF INVESTIGATION

- **Consequences of implantation and mixing are poorly characterized in modeling of plasma etching.**
- **Usually only included in compute-intensive molecular dynamics simulations.**
- **Incorporate implantation, mixing, and sputtering into Monte-Carlo Feature Profile Model coupled to equipment scale plasma models.**
- **Investigate:**
 - **Scaling of implantation, mixing, and etching selectivity.**
 - **Degradation and cross-linking of PR surface due to energetic bombardment.**
- **Goals**
 - **Characterize mixing damage during etching.**
 - **Develop strategies to preserve pattern transferring while minimizing damage.**

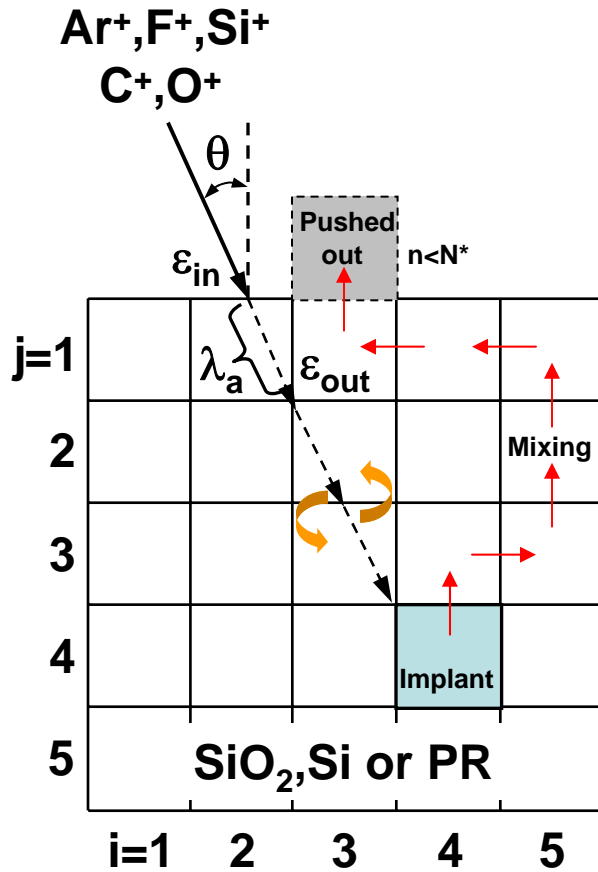
DESCRIPTION OF MODEL



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IMPLANTATION MODEL



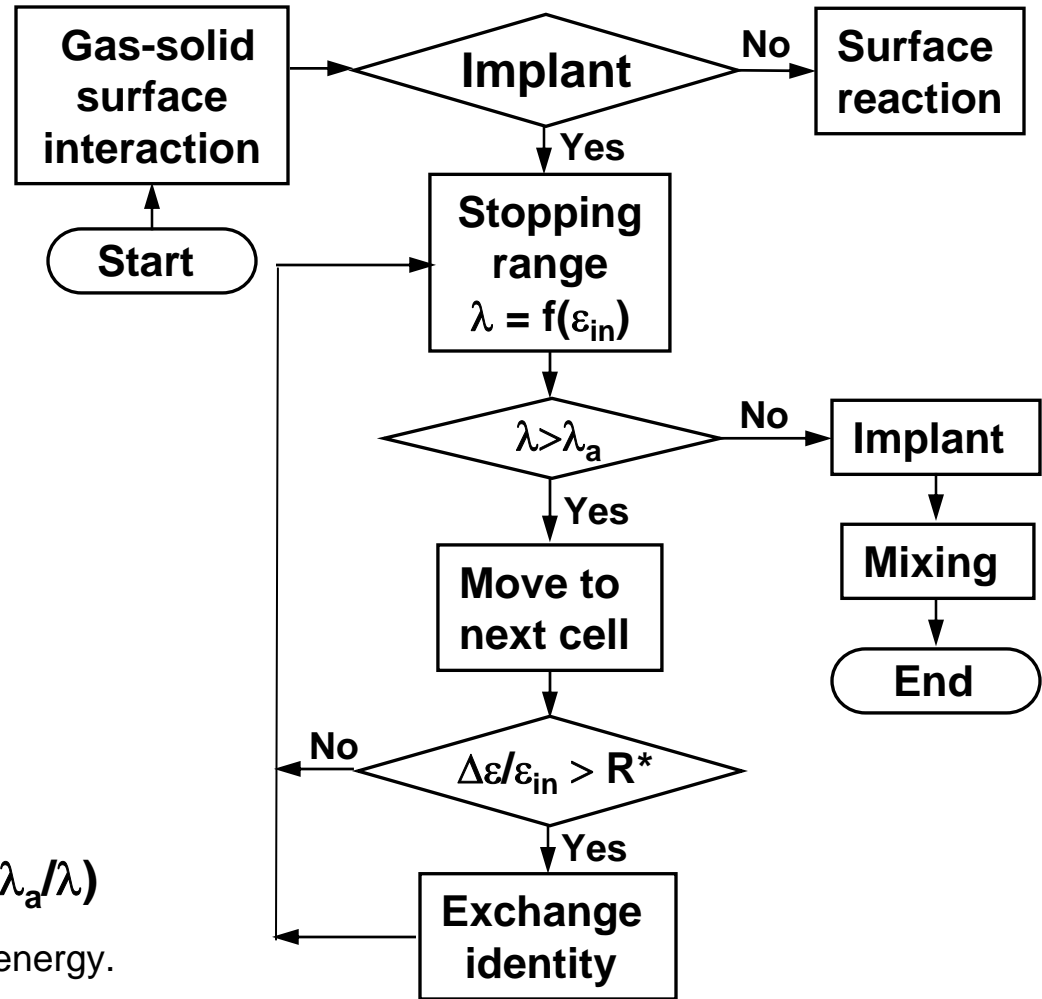
Within one cell: $\epsilon_{out} = \epsilon_{in} \exp(-\lambda_a/\lambda)$

Where ϵ_{in} = incident energy; ϵ_{out} = left energy.

λ_a = Actual length that the particle travels.

λ = Calculated stopping range $f(\epsilon_{in})$.

*n = mixing step; N = allowed maximum mixing step.



*R = Random number

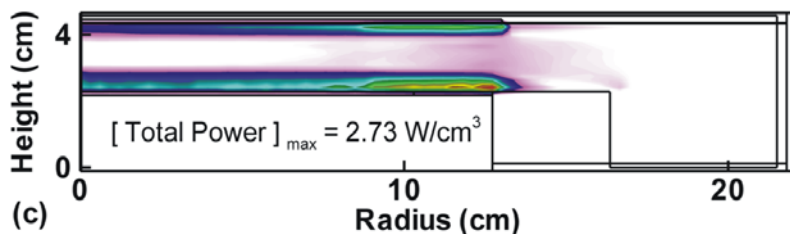
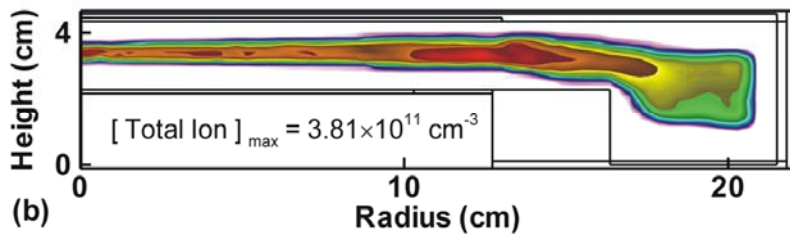
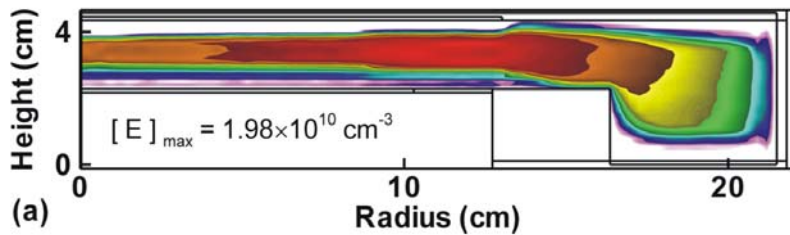
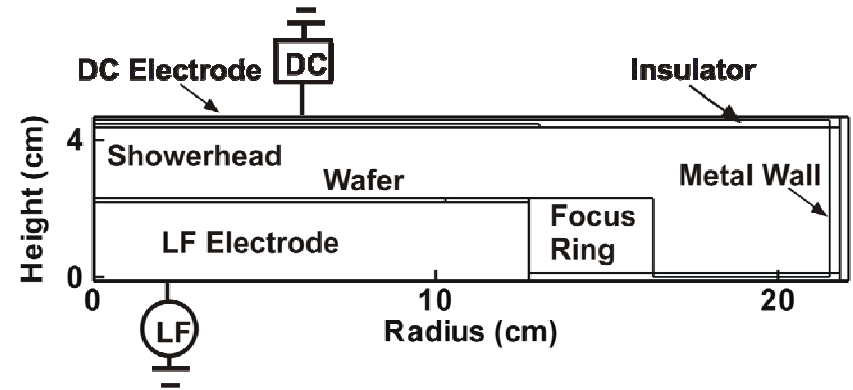
SURFACE REACTION MECHANISM

- Etching of SiO_2 is dominantly through a formation of a fluorocarbon complex.
 - $\text{SiO}_2(\text{s}) + \text{C}_x\text{F}_y^+(\text{g}) \rightarrow \text{SiO}_2^*(\text{s}) + \text{C}_x\text{F}_y(\text{g})$
 - $\text{SiO}_2^*(\text{s}) + \text{C}_x\text{F}_y(\text{g}) \rightarrow \text{SiO}_2\text{C}_x\text{F}_y(\text{s})$
 - $\text{SiO}_2\text{C}_x\text{F}_y(\text{s}) + \text{C}_x\text{F}_y^+(\text{g}) \rightarrow \text{SiF}_y(\text{g}) + \text{CO}_2(\text{g}) + \text{C}_x\text{F}_y(\text{g})$
 - Further deposition by $\text{C}_x\text{F}_y(\text{g})$ produces thicker polymer layers.
- Example reaction of surface dissociation.
 - $\text{M}(\text{s}) + \text{C}_x\text{F}_y^+(\text{g}) \rightarrow \text{M}(\text{s}) + \text{C}_{x-1}\text{F}_{y-1}(\text{g}) + \text{C}(\text{g}) + \text{F}(\text{g})$
- Sputtering of PR and redeposition.
 - $\text{PR}(\text{s}) + \text{Ar}^+(\text{g}) \rightarrow \text{PR}_2(\text{s}) + \text{Ar}(\text{g}) + \text{H}(\text{g}) + \text{O}(\text{g})$
 - $\text{PR}(\text{s}) + \text{C}_x\text{F}_y^+(\text{g}) \rightarrow \text{PR}(\text{s}) + \text{C}_x\text{F}_y(\text{g})$
 - $\text{PR}(\text{g}) + \text{SiO}_2\text{C}_x\text{F}_y(\text{s}) \rightarrow \text{SiO}_2\text{C}_x\text{F}_y(\text{s}) + \text{PR}(\text{s})$

*PR2 = cross-linked PR

FLUOROCARBON ETCHING OF SiO₂

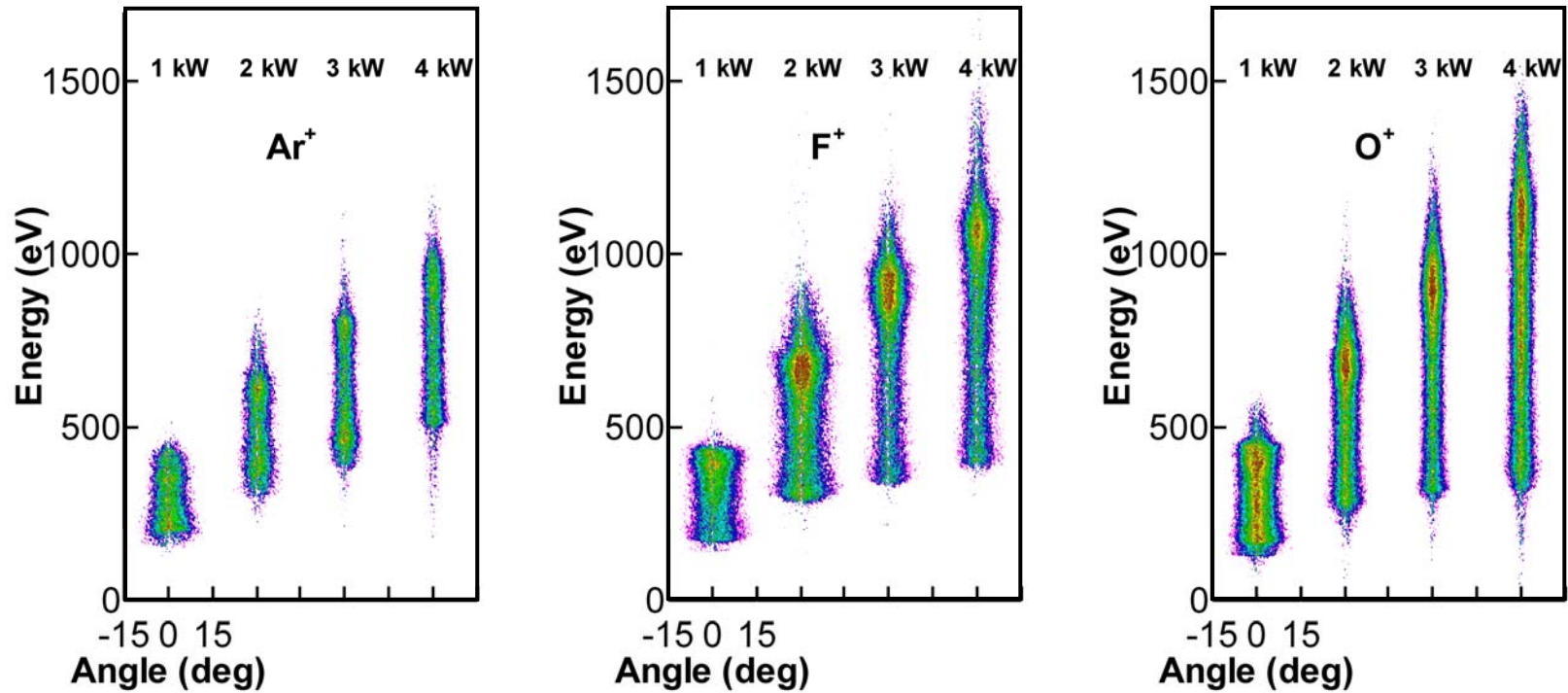
- DC augmented single frequency capacitively coupled plasma (CCP) reactor.
- DC: Top electrode RF: Substrate



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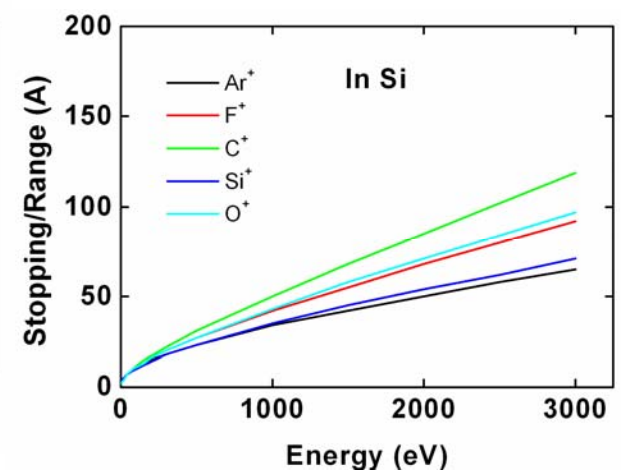
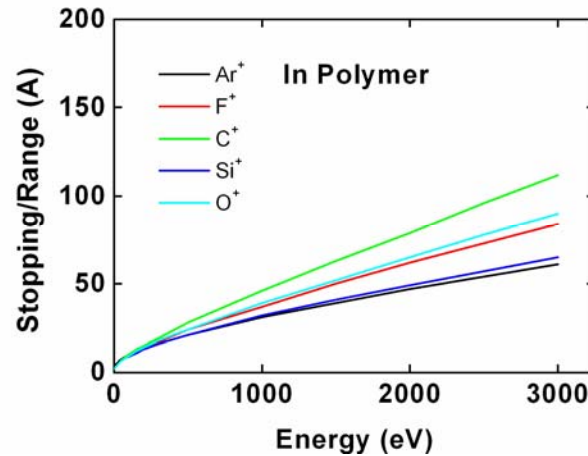
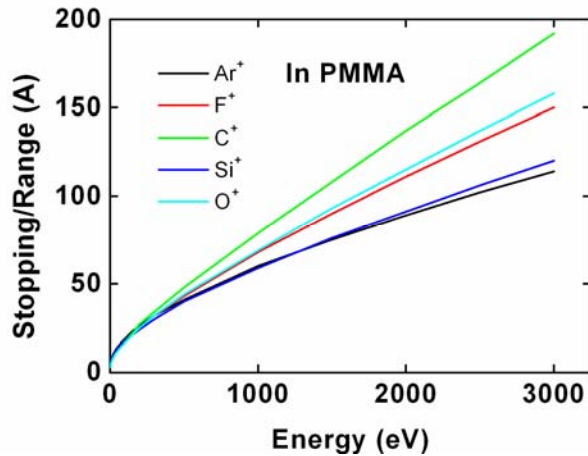
- Plasma tends to be edge peaked due to electric field enhancement.
- Plasma densities in excess of 10^{11} cm^{-3} .
- Ar/C₄F₈/O₂ = 80/15/5, 300 sccm, 40 mTorr, RF 1 kW at 10 MHz, DC 200 W/-250 V.

ION ENERGY ANGULAR DISTRIBUTIONS (IEADs)



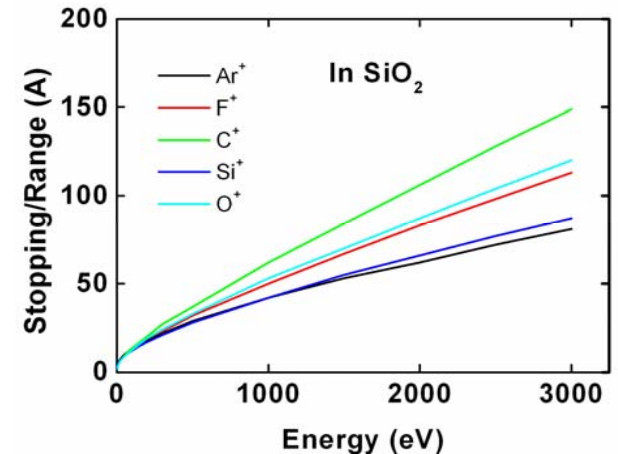
- Peak of ion energy ranges from 300 to 1200 eV for 1 – 4 kW bias power.
- Angle distribution spreads from -10 to 10 degree .
- Stopping range ranges from 0 to 70 Å.

STOPPING RANGE IN PR, POLYMER, Si, AND SiO₂

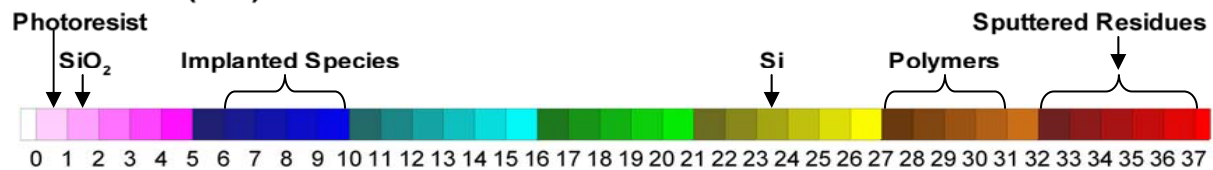
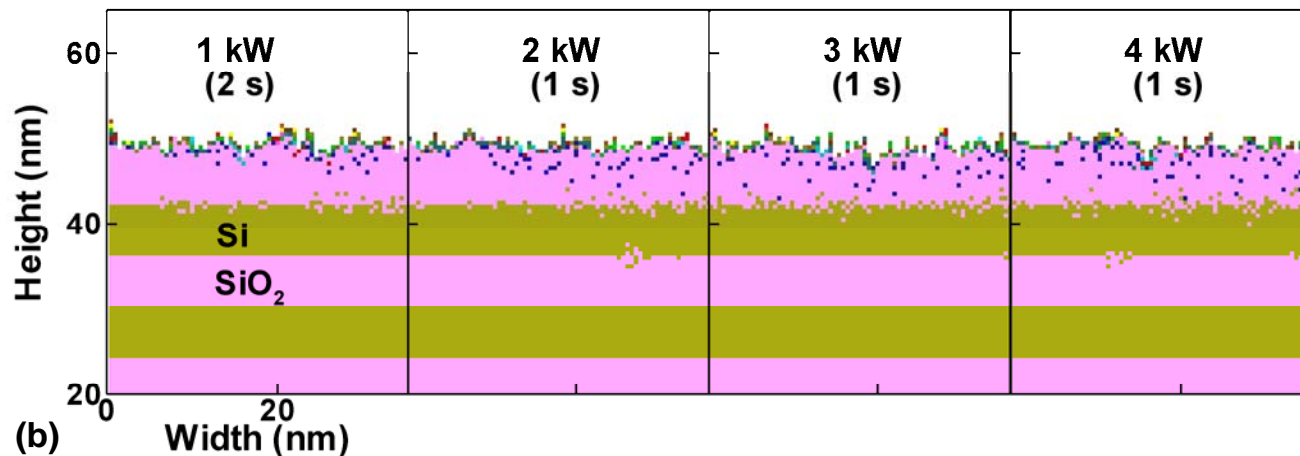
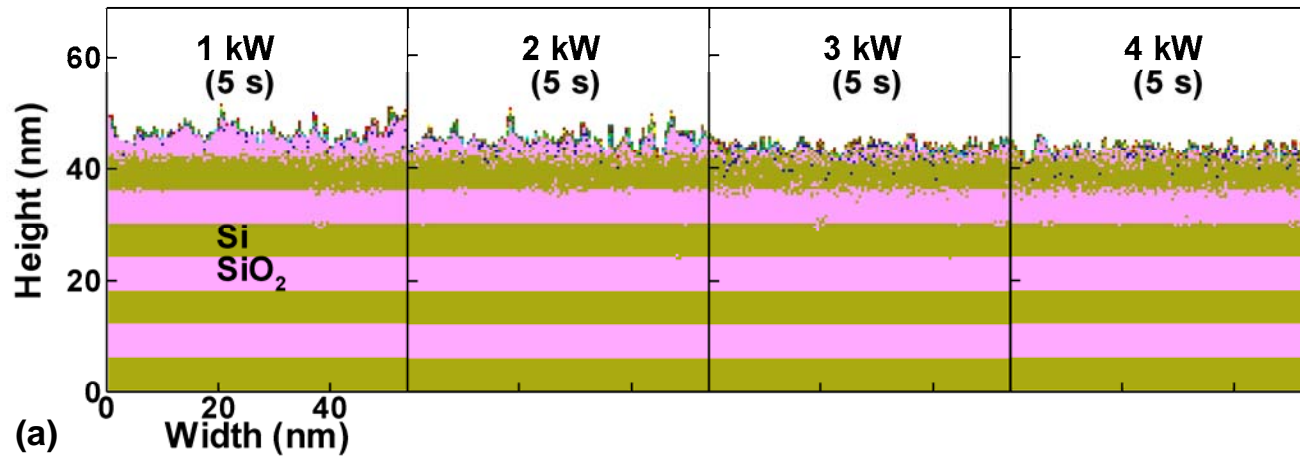


- Stopping range increases with increasing energy.
- At specific energy, implanting depth:
 $PR(\text{PMMA}^*) > \text{SiO}_2 > \text{Si} \geq \text{Polymer}$
 *PMMA = Polymethylmethacrylate
- Data from SRIM (the Stopping and Range of Ion in Matter)

$$\text{Stopping range } \lambda = f(\epsilon_{\text{incident}})$$

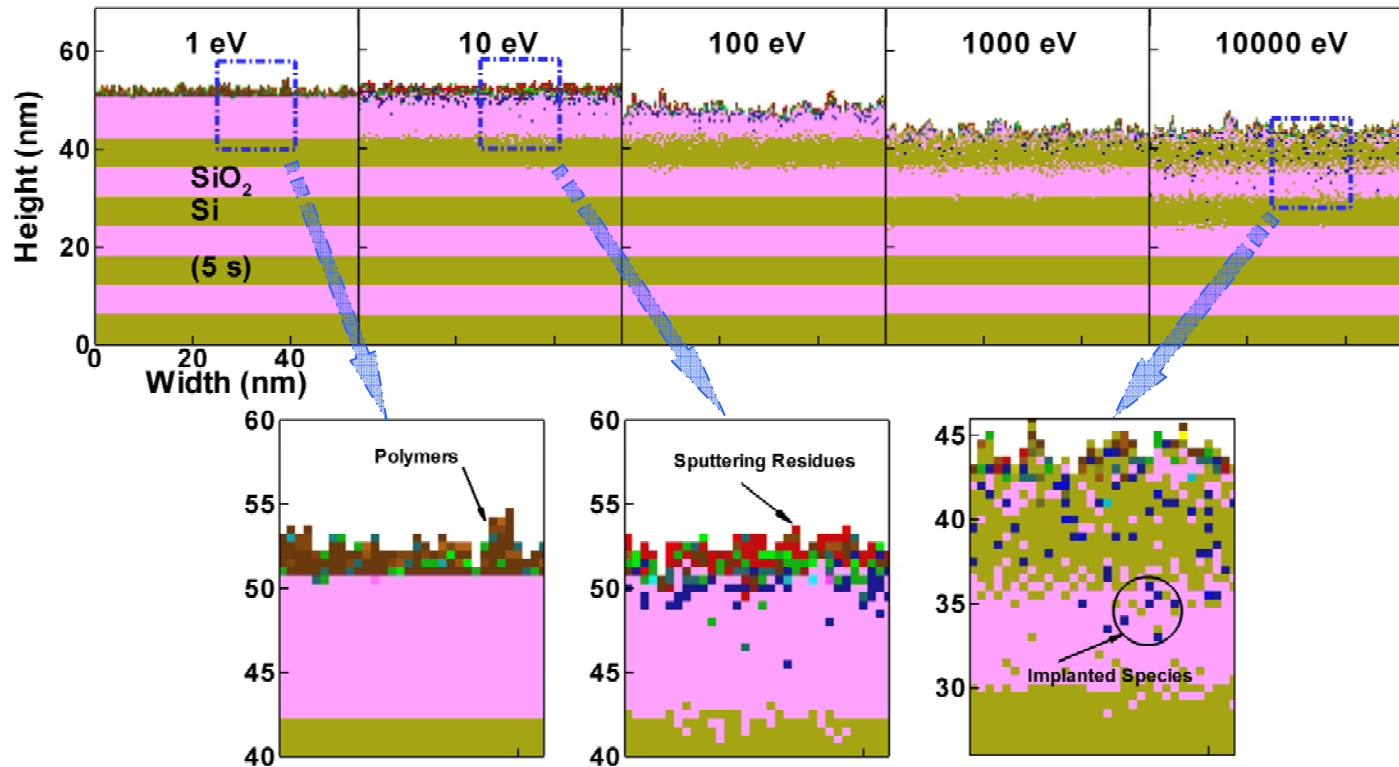


IMPLANTING AND MIXING DEPTH vs BIAS POWER



- After 5 s
- Etch rates, degree of mixing and depth of implantation increase with bias power.
- After same etch level

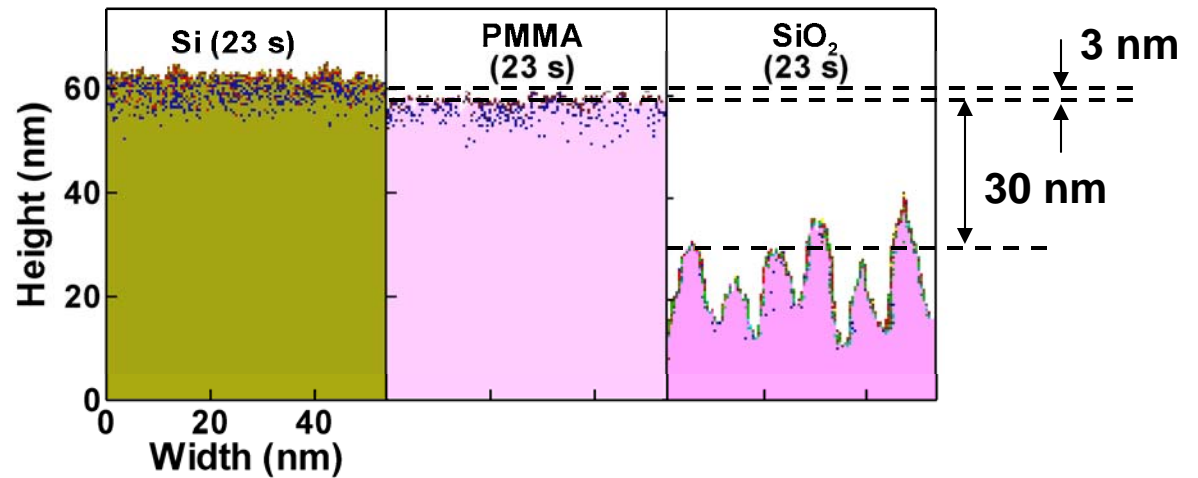
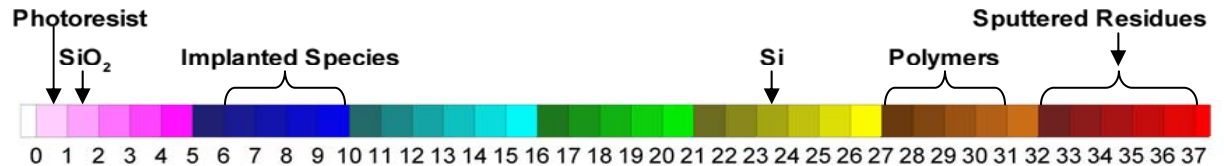
IMPLANTING AND MIXING DEPTH vs ENERGY



- Only polymer deposition occurs at 1 eV.
- Sputtering, implanting and deposition coexist at 10 eV.
- Depth of implantation and mixing increases with increasing ion energy (100 eV~10 keV).

ETCHING SELECTIVITY

- Etch stop occurs at Si surface due to low reaction rate with CF_x .
- Etching selectivity for SiO_2 /PR(PMMA) is around 10.
- The roughness of SiO_2 surface is due to non-uniform polymer deposition.



Micro-masking \longrightarrow Micro-trenching

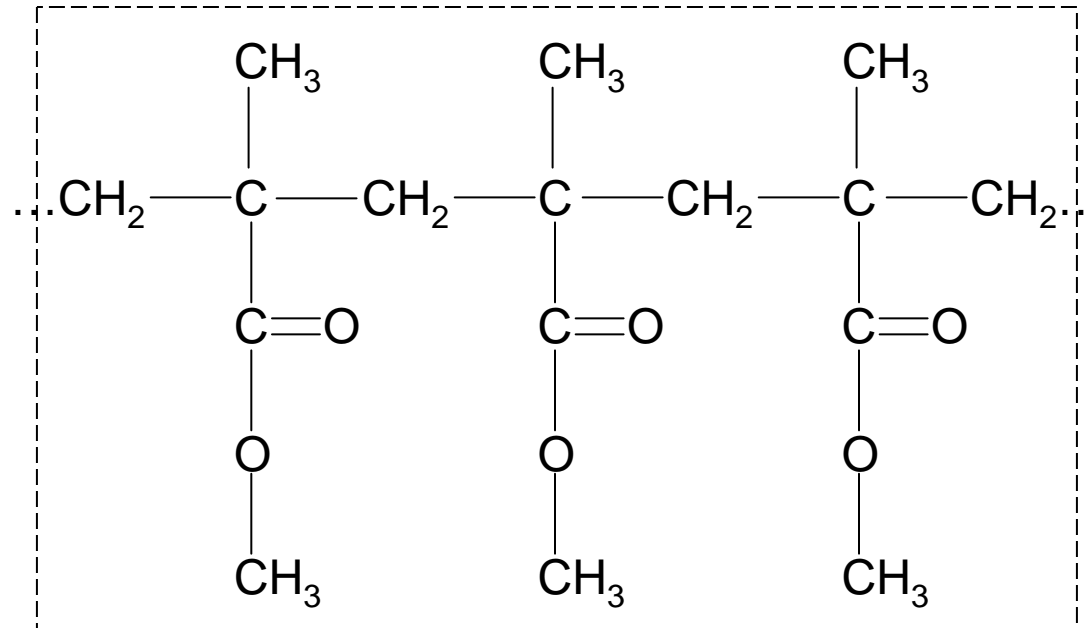
- $Ar/C_4F_8/O_2 = 80/15/5$, 4 kW, 300sccm, 40mTorr, DC 200W/-250V.

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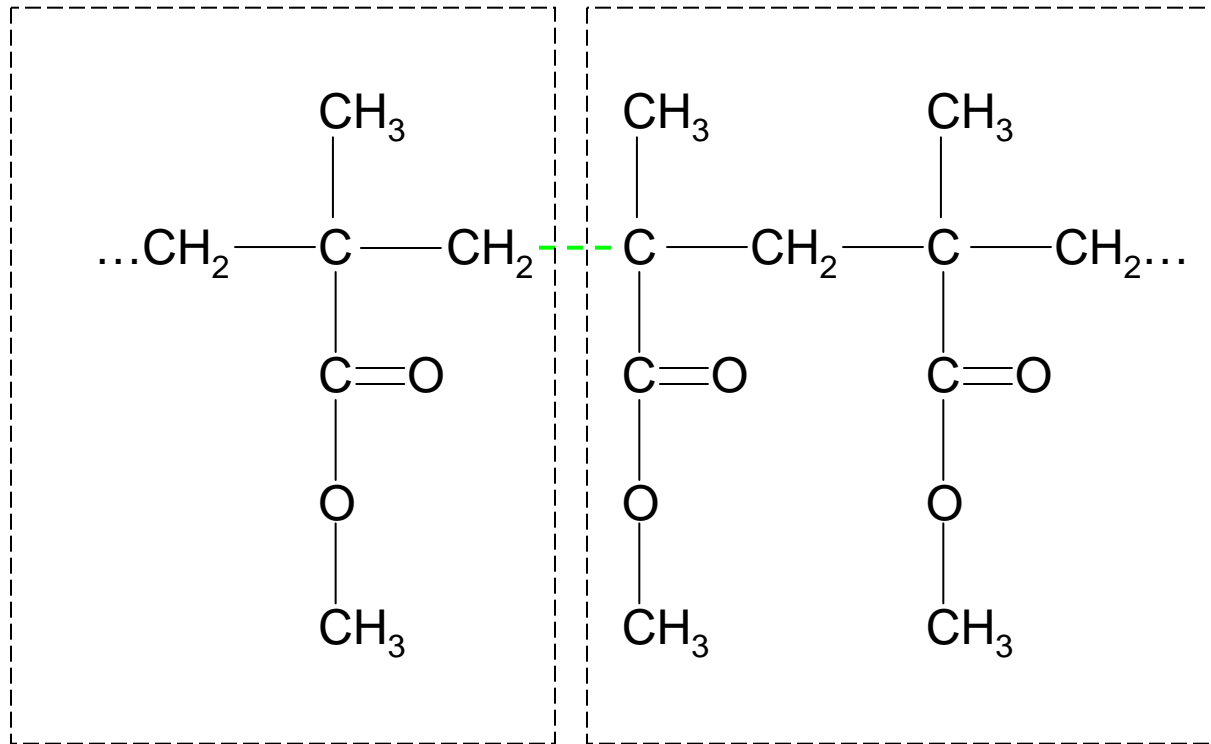
MECHANISM FOR DEGRADATION AND CROSS-LINKING OF PR

Structure of PR(PMMA)



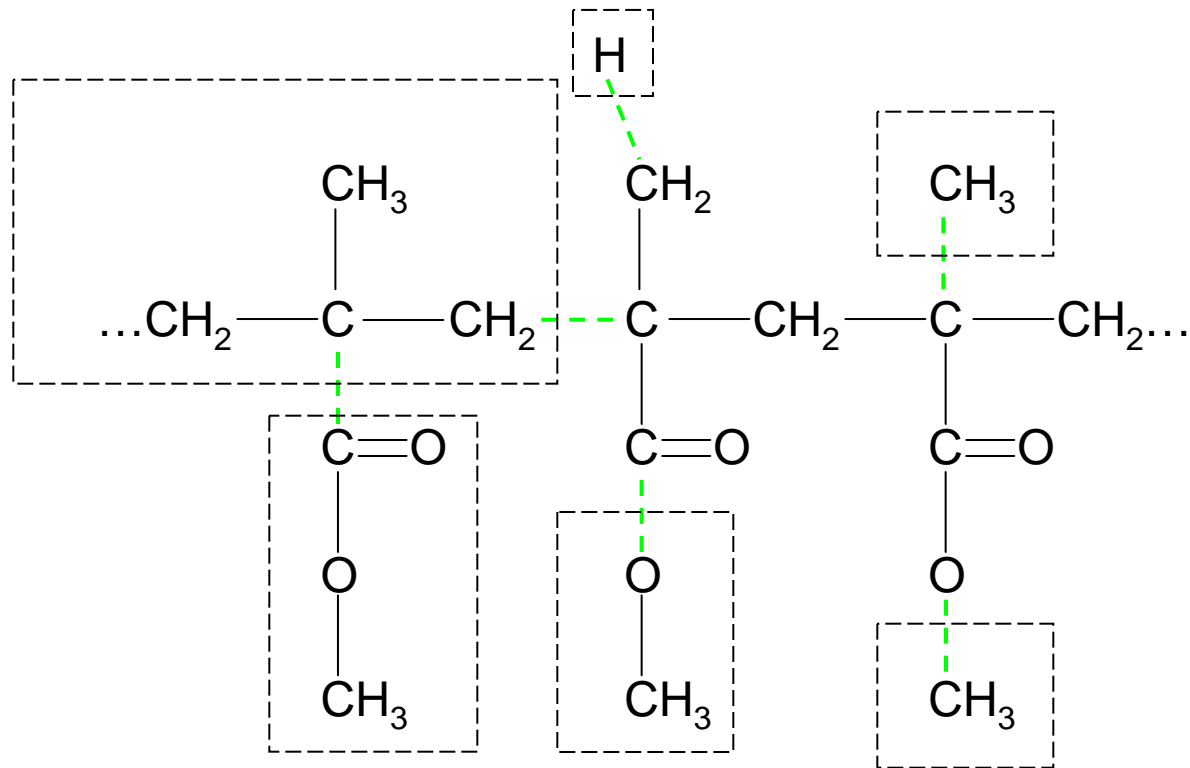
- PR molecule is degraded due to energetic ion sputtering.
- Degraded PR segments are more easily eroded.
- Newly formed dangling bonds are recombined (cross-linking), and cross-linked PR forms a “hard crust”.
- Sputtering yield is calculated using SRIM.

DEGRADATION OF PR (SPUTTERING)



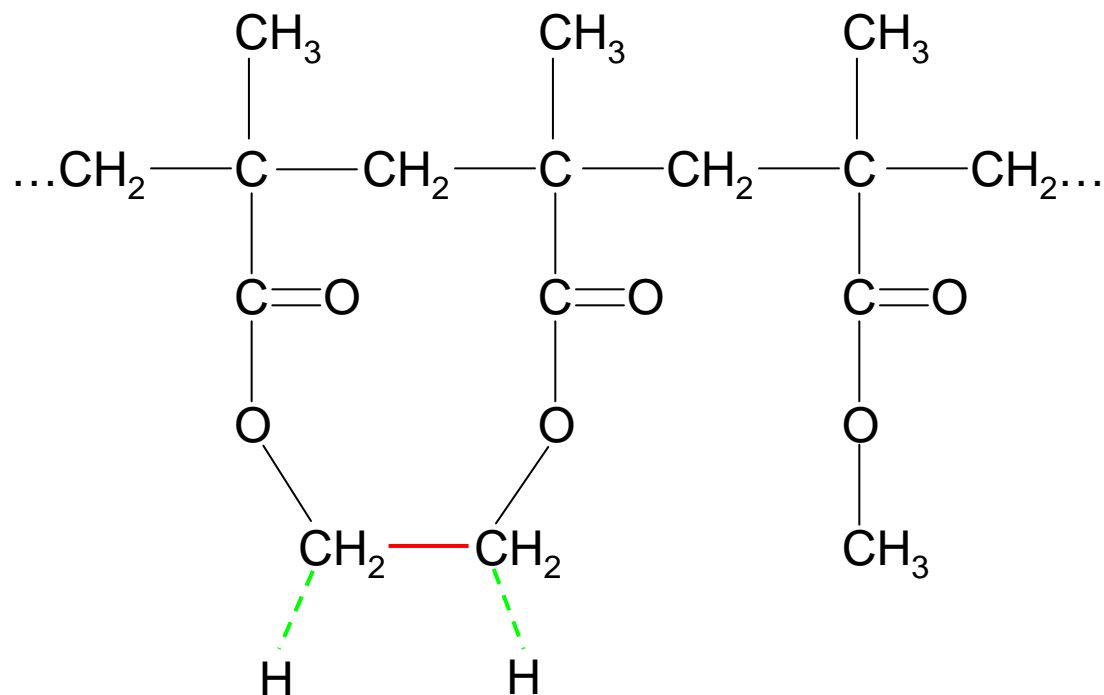
----- Broken Bond

DEGRADATION OF PR (SPUTTERING)



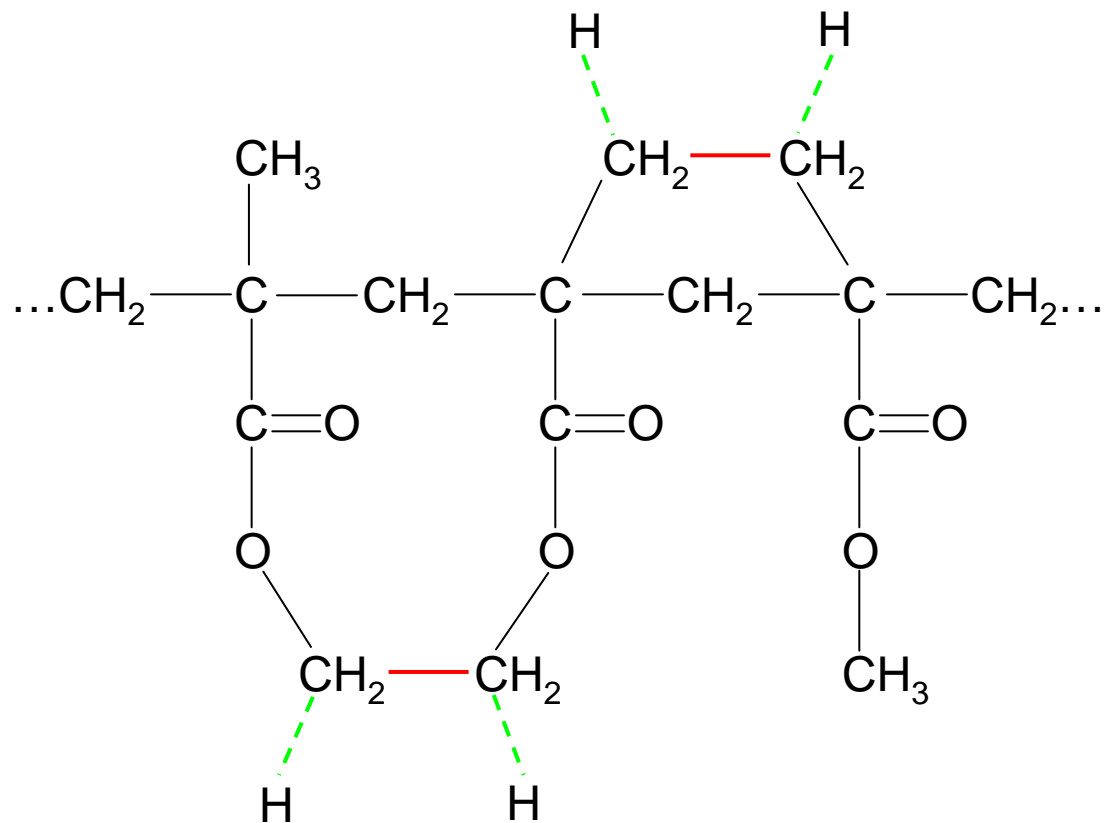
----- Broken Bond

INTER-MOLECULE CROSS-LINKING



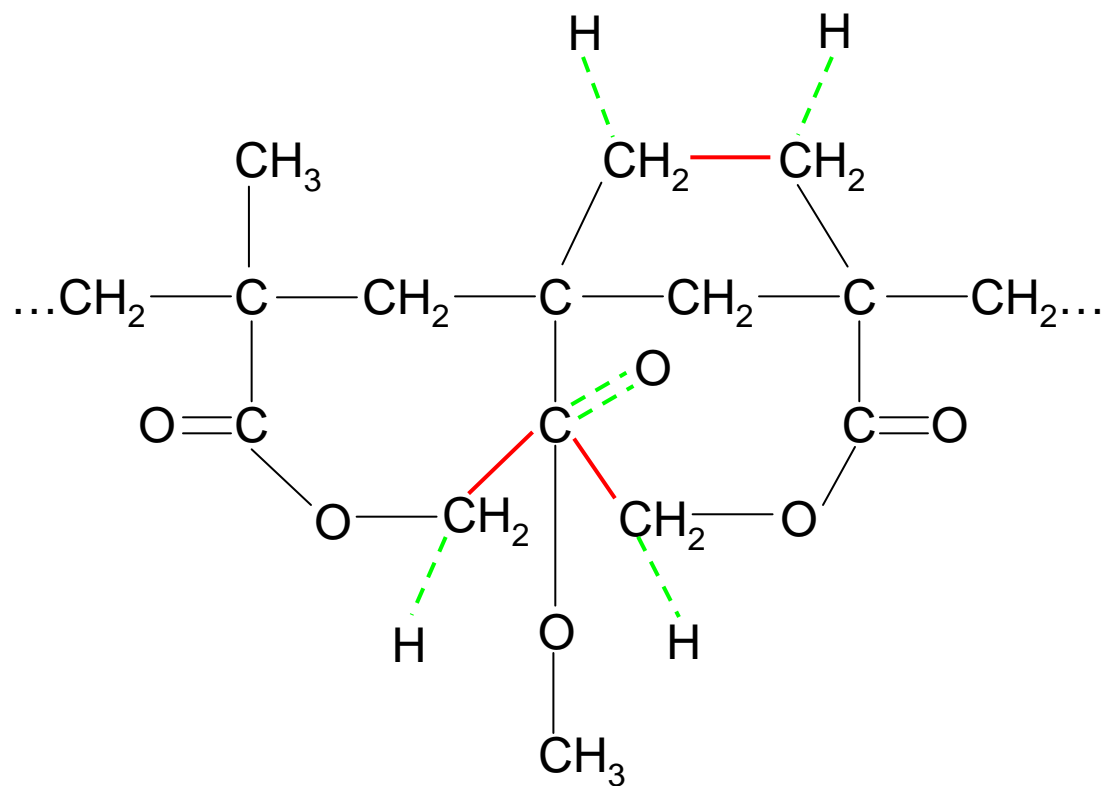
----- Broken Bond
----- New Bond

INTER-MOLECULE CROSS-LINKING



----- Broken Bond
----- New Bond

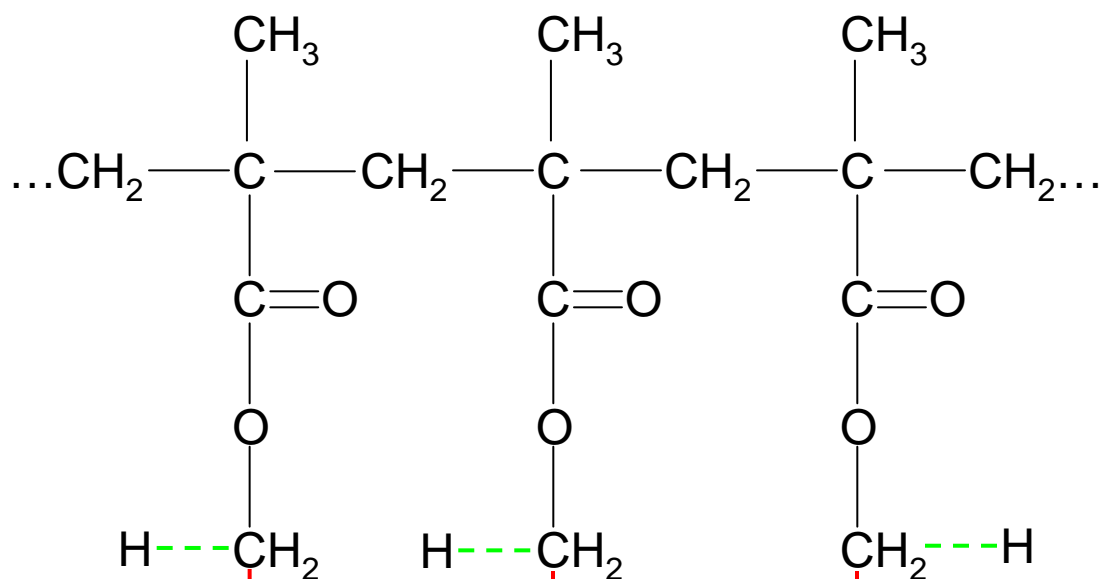
INTER-MOLECULE CROSS-LINKING



----- Broken Bond
----- New Bond

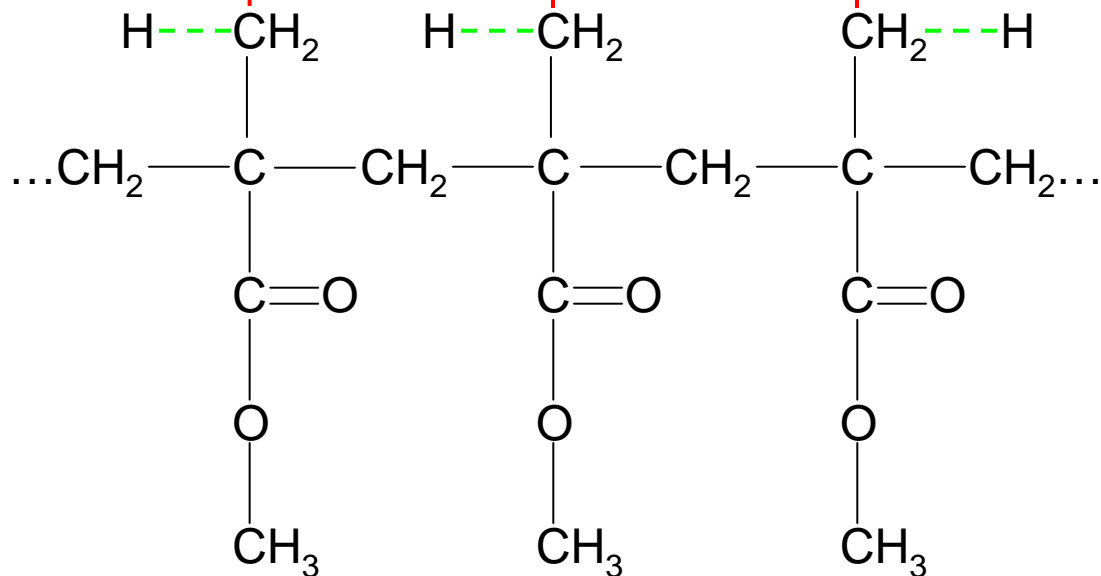
INTRA-MOLECULE CROSS-LINKING

Molecule A



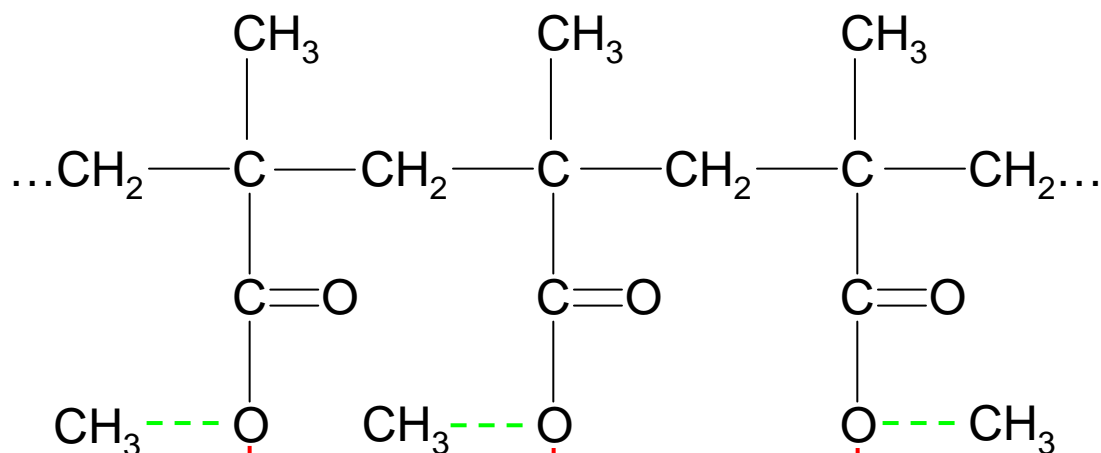
----- Broken Bond
----- New Bond

Molecule B



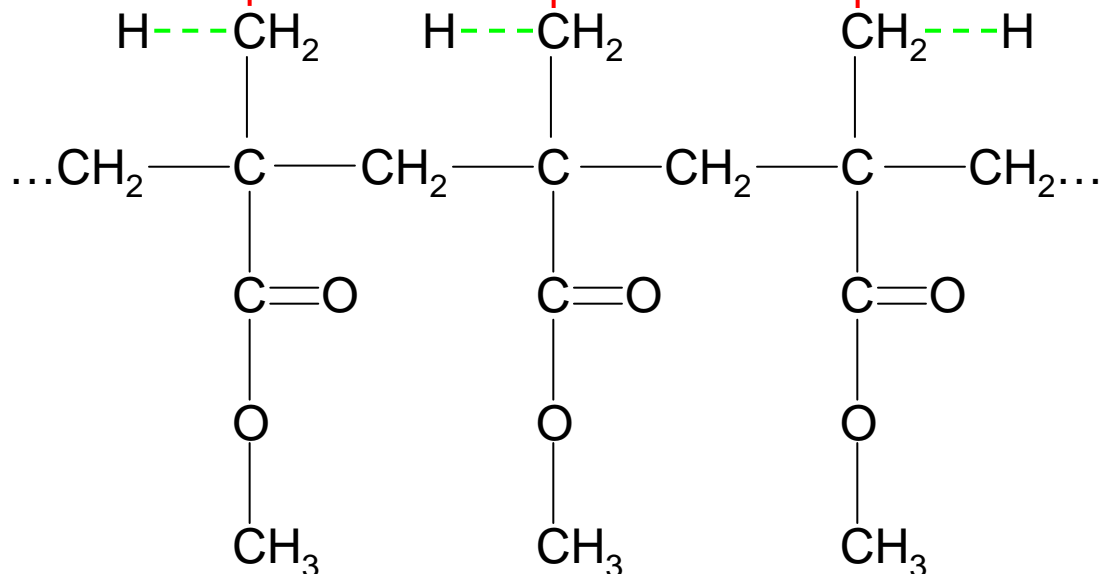
INTRA-MOLECULE CROSS-LINKING

Molecule A

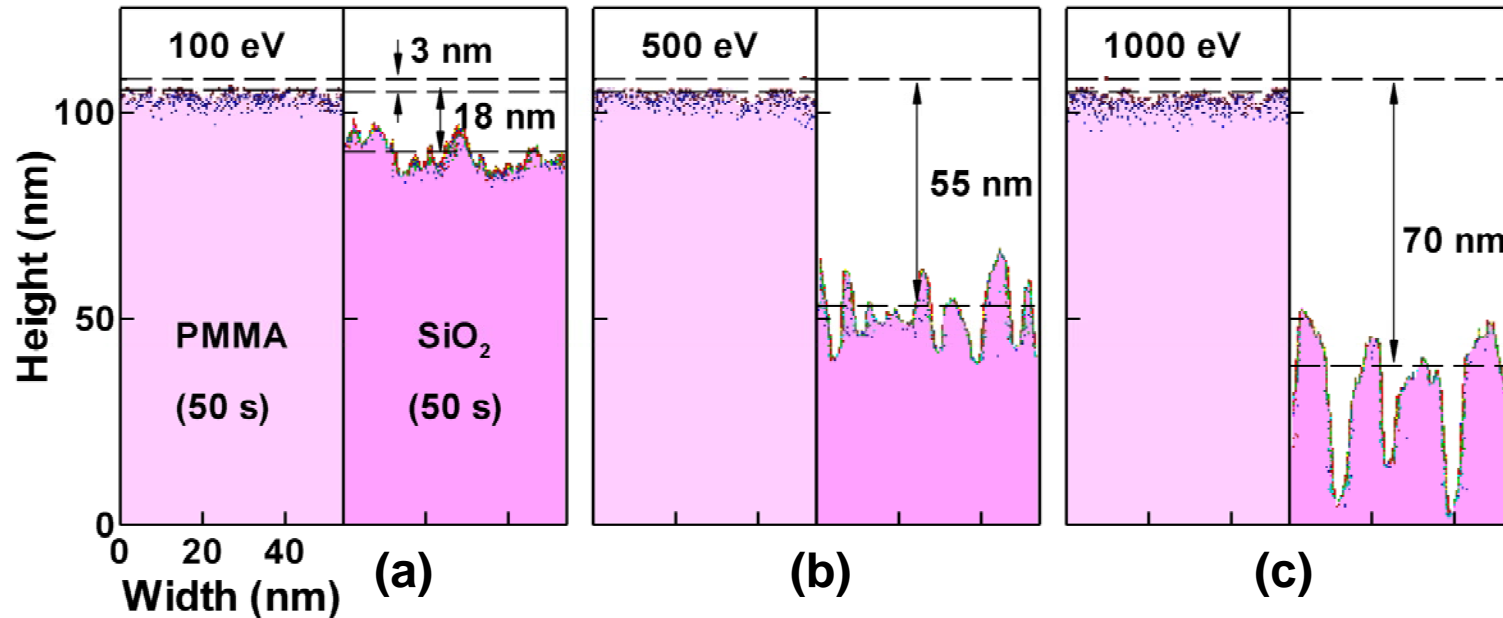


----- Broken Bond
----- New Bond

Molecule B



ETCHING SELECTIVITY vs ENERGY

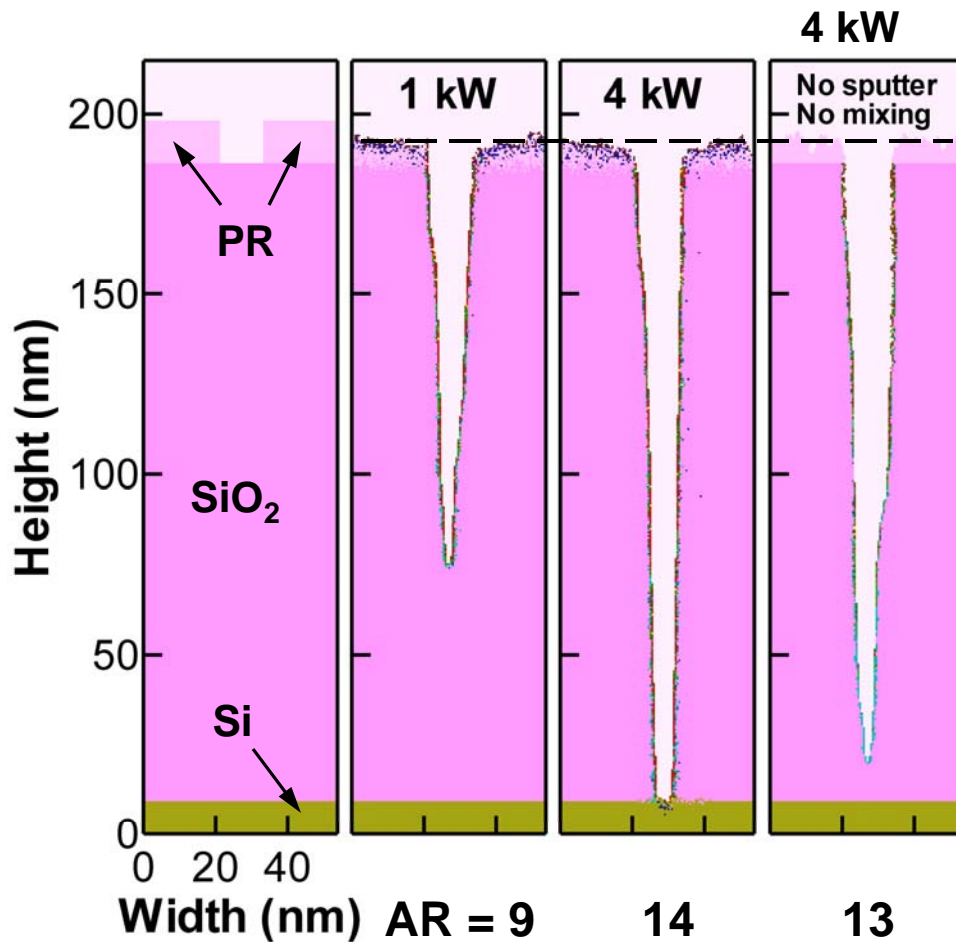


- Etching rate for SiO₂ increases with increasing ion energy.
- Balance between sputtering and cross-linking (more resistive to etching) on PR(PMMA) surface results in similar etching rate for all energies.
- Surface roughness of SiO₂ increases as etching proceeds due to micro-masking.
- Etching selectivity (SiO₂/PR): 100 eV, 6; 500 eV, 18; 1000 eV, 23.

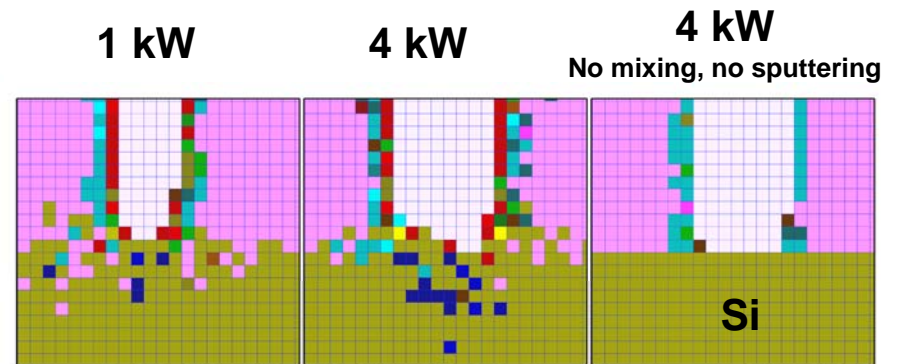
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ETCHING SELECTIVITY vs RF BIAS POWER



- Ar/C₄F₈/O₂ = 80/15/5, 300 sccm, 40 mTorr, 10 MHz, DC 200 W/-250 V.



- At similar etching level of PR, aspect ratio (AR) of the trench increases with bias power.
- Etching selectivity increases when PR is cross-linked.
- Si is damaged during over-etch by implantation and mixing.

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CONCLUDING REMARKS

- Algorithms have been incorporated into the MCFPM to predict implanting and mixing.
 - Depth of implanting and mixing increases with increasing bias power and ion energy.
 - More damage is obtained at higher etching rate.
- PR surface is cross-linked due to sputtering.
 - At higher bias power and ion energy, etching selectivity for SiO₂/PR is better.
- Strategies to be developed for high power processing:
 - Protect pattern transferring at higher etching rate.
 - Reduce damage during over-etch.