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CHE323/CHE384
 Chemical Processes for Micro- and Nanofabrication
www.lithoguru.com/scientist/CHE323

Lecture 24 Chemical Vapor Deposition, part 1

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Reading:
 Chapter 13, *Fabrication Engineering at the Micro- and Nanoscale*, 4th edition, Campbell

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Chemical Vapor Deposition

- CVD is used to deposit most dielectrics and silicon, but also for some metals
- Performed in furnaces, like oxidation and diffusion, and in newer single-wafer reactors
- Chemical reactions produce the desired species being deposited

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Chemical Vapor Deposition

- Gas-phase (homogeneous) reactions
 - Reactions occur in the gas phase
 - Solid is produced, that then lands on the wafer
 - Poor uniformity, many particles
- Solid surface (heterogeneous) reactions
 - Reaction occurs on the wafer surface
 - Solid formed is the deposited film
 - This is what we try to achieve

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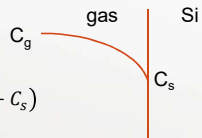
Simplified CVD Mechanism

- Transport of reactants to wafer surface (diffusion)
- Adsorption of reactants on surface
- Reaction
 - Assumed to be rate-limiting step
- Desorption of by-products
- Transport of by-products into gas stream

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CVD Mechanism, Step 1



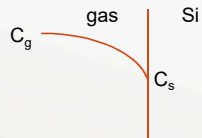
$$J_1 = D \frac{dC}{dx} \approx \frac{D_g}{\delta} (C_g - C_s) = h_g (C_g - C_s)$$

J_1 = flux of reactant to wafer surface
 D_g = diffusivity of reactant in gas
 C_g = bulk reactant concentration = $\frac{n}{V} = \frac{P_g}{kT}$
 C_s = surface gas concentration
 δ = boundary layer thickness
 $h_g = D_g/\delta$ = mass transfer coefficient

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CVD Mechanism, Step 2



$$J_2 = k_s C_s$$

J_2 = flux of reactant as it reacts
 k_s = reaction rate constant
 C_s = reactant concentration at wafer surface

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Steady State

- Assume that the surface reaction is the rate-limiting step
- Steady state is reached where all fluxes are equal

$$J_1 = J_2 = J_{SS}$$

- Solve for the unknown concentration (C_S) and eliminate from equation

$$C_S = \frac{h_g C_g}{h_g + k_s}$$

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Simplified Rate Equation

- Steady State reaction rate

$$J_{SS} = k_T C_g \quad k_T = \frac{h_g k_s}{h_g + k_s}$$

- For $N = \# \text{ atoms/cm}^3$ in deposited film,

$$\text{deposition rate} = v = \frac{J_{SS}}{N}$$

- Overall deposition rate

$$v = \frac{k_T}{N} C_g = \frac{k_T}{N} \left(\frac{1}{kT} \right) P_g$$

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Simplified Rate Equation

- Deposition rate

$$v = \frac{k_T}{N} \left(\frac{1}{kT} \right) P_g \quad k_T = \frac{h_g k_s}{h_g + k_s}$$

- Two regimes,

if $k_s \gg h_g, \quad k_T \approx h_g$ (diffusion - controlled)

if $k_s \ll h_g, \quad k_T \approx k_s$ (reaction - controlled)

Note: P_g must be kept low to prevent homogeneous reaction

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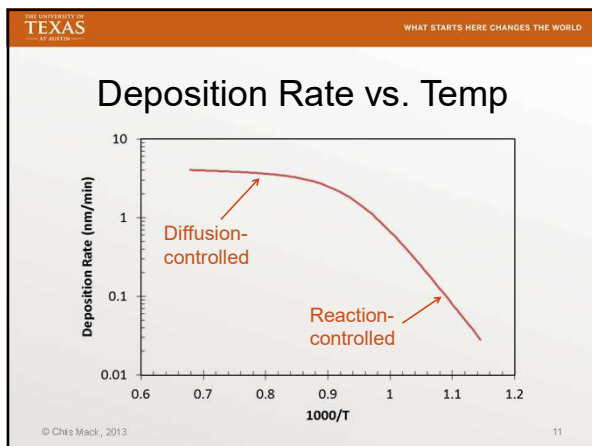
Effect of Temperature

- In general, $E_{a(\text{reaction})} \gg E_{a(\text{gas diffusion})}$

$$D_g \propto T^{\frac{3}{2}} \frac{P_g}{P_T}, \quad \delta \text{ varies slowly with } T$$

- At high temperatures, $k_s \gg h_g$
 - Diffusion-limited regime
- At low temperatures, $k_s \ll h_g$
 - Reaction-limited regime

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Lecture 24: What have we learned?

- Why is heterogeneous CVD preferred over homogeneous CVD?
- What are the two steps in our very simplified mechanism for CVD?
- Explain diffusion-controlled vs. reaction-controlled regimes for CVD?

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