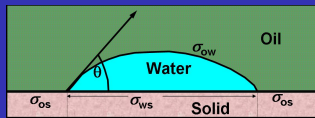


Multiphase Flow and Heat Transfer



Wettability

Sudheer Siddapureddy

sudheer@iitp.ac.in



Department of Mechanical Engineering
Indian Institution of Technology Patna



- Usually, a liquid-vapor phase change is accomplished by transferring energy through the walls of a container or channel into or out of a two-phase system.
- The vaporization or condensation process ultimately takes place at the liquid-vapor interface.
- However, the contact through which the energy is transferred will strongly affect the resulting heat and mass transfer in the system.
- The performance of heat transfer equipment in which vaporization or condensation occurs may depend strongly on the way that the two phases contact the solid walls.



- Liquids with weak affinities for a solid wall will collect themselves into beads while those with high affinities for solid will form film to maximize the liquid-solid contact area.
- The affinity of liquids for solids - **wettability of the fluid.**



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wetting
 $\Theta = 0^\circ$

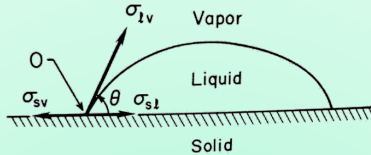


partial wetting
 $0^\circ < \Theta < 180^\circ$



Nonwetting
 $\Theta = 180^\circ$

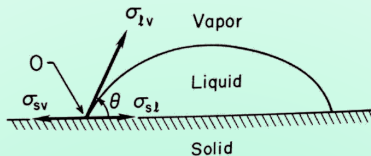
Neumann's Formula or Young's Equation



$$\sigma_{sv} = \sigma_{sl} + \sigma_{lv} \cos \theta$$

Vertical force $\sigma_{lv} \sin \theta$ must be balanced by a vertical reaction force in the solid.

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Vertical force $\sigma_{lv} \sin \theta$ must be balanced by a vertical reaction force in the solid.

- Small and also modulus of elasticity of solid is high
- No deformation occurs

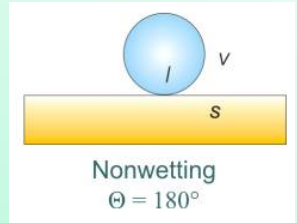
σ_{sl} and σ_{sv} are not available easily.

Neumann's Formula or Young's Equation



As $\theta \rightarrow 180^\circ$ (if g is negligible)

- Liquid droplet - sphere
- One point of contact on solid
- Completely non-wetting

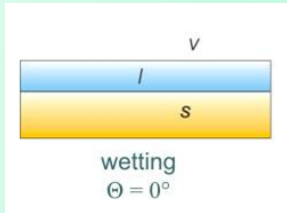
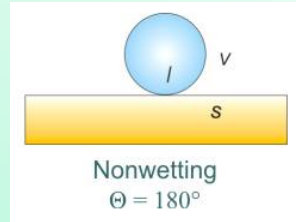


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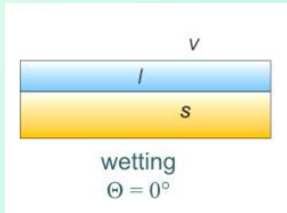
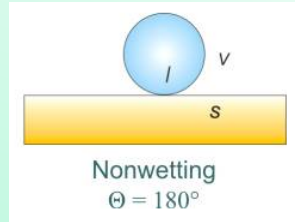
- A thin film configuration
- Completely wet the solid

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As $\theta \rightarrow 0^\circ$

- A thin film configuration
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- Wetting liquid: As $0^\circ < \theta < 90^\circ$
- Non-wetting liquid: As $90^\circ < \theta < 180^\circ$
one point of contact on solid

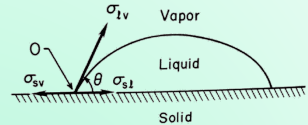
Neumann's Formula or Young's Equation



$$\sigma_{sv} = \sigma_{sl} + \sigma_{lv} \cos \theta; \quad |\cos \theta| \neq 1$$

At equilibrium:

$$\left| \frac{\sigma_{sv} - \sigma_{sl}}{\sigma_{lv}} \right| < 1$$



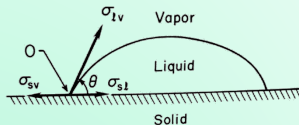
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Never happens for a droplet surrounded by its vapor, but could happen for a liquid droplet in another immiscible liquid.

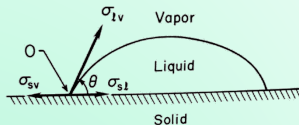
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Never happens for a droplet surrounded by its vapor, but could happen for a liquid droplet in another immiscible liquid.

If $\frac{\sigma_{sv} - \sigma_{sl}}{\sigma_{lv}} > 1$, σ_{sv} pulls the contact line, $\theta \rightarrow 0^\circ$

Until the film becomes so thin that molecular interactions come into play.



$$Sp_{ls} = \sigma_{sv} - (\sigma_{lv} + \sigma_{sl})$$

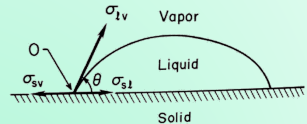
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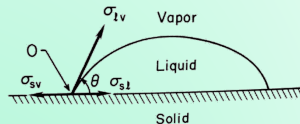
$$Sp_{ls} = \sigma_{lv} (\cos \theta - 1)$$



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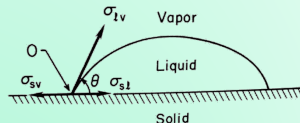
$Sp_{ls} > 0$, the liquid will wet the solid and spontaneously spread into a thin film.

$Sp_{ls} < 0$, the liquid will partially wet the solid and establish an equilibrium contact angle.

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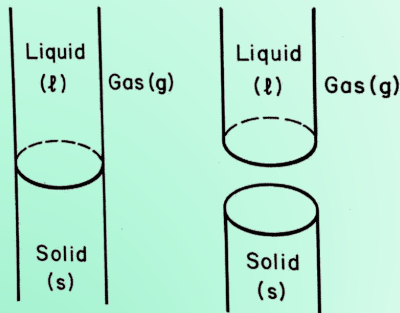
These results are theoretical and no lack of σ_{sv} and σ_{sl} .

Cylindrical column formed by l , s and low density gas, g . The net reversible work required per unit of interface area:

$$w_{sl} = \sigma_{lg} + \sigma_{sg} - \sigma_{sl}$$

Essentially,
Forming 2 new interfaces
Breaking one interface

Adhesion: due to the minimum reversible work required to tear the liquid off the solid surface.





Now consider, instead of solid-liquid column there is only one liquid column. To tear a single liquid column in half:

$$w_{ll} = 2\sigma_{lg}$$

Essentially, two interfaces are formed without breaking any interface.

Cohesion: work required to break internal bonds of the material.



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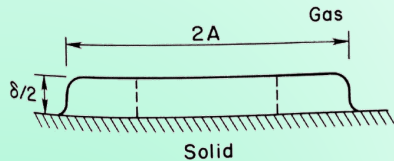
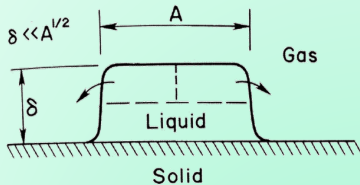
Adhesion - for dissimilar particles

Cohesion - for similar particles

Work per Unit Area Required for Spreading



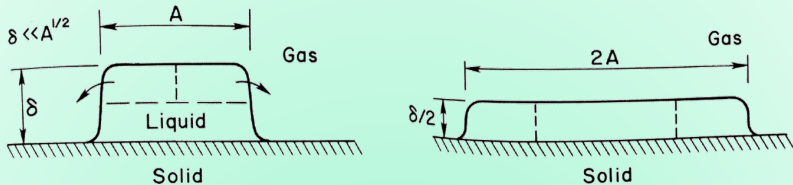
Now let's march further. Let's say there is a liquid column of area A , and height δ . We are making it into a liquid column of area $2A$ and height of $\delta/2$.



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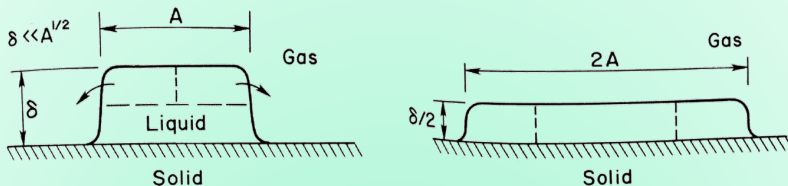


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$$w_{sp} = \sigma_{lg} + \sigma_{sl} - \sigma_{sg}$$

$$= -Sp_{ls}$$

Work interaction is negative (Sp_{ls} is positive for spreading). Work could be extracted if we can ($= Sp_{ls}$).



$$Sp_{ls} = -w_{sp} = w_{sl} - w_{ll}$$

Spreading coefficient = difference between $\left\{ \begin{array}{l} \text{the work of adhesion} \\ \text{the work of cohesion} \end{array} \right.$

Sp_{ls} indicates the tendency of the liquid to adhere to the solid relative to its internal cohesive forces.



$$w_{sl} = \sigma_{lg} + \sigma_{sg} - \sigma_{sl}$$

Applying it to any two solid or liquid phases a and b , and a low density gas or vapor phase g ,

$$w_{ab} = \sigma_{bg} + \sigma_{ag} - \sigma_{ab}$$





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$$\sigma_{ab} = \sigma_{bg} + \sigma_{ag} - w_{ab}$$

The work of adhesion is approximately given as:

$$w_{ab} \cong 2 (\sigma_{ag}\sigma_{bg})^{\frac{1}{2}}$$

$$\sigma_{ab} = \sigma_{bg} + \sigma_{ag} - 2 (\sigma_{ag}\sigma_{bg})^{\frac{1}{2}}$$



For water and hexane in contact with air at 20°C, σ_{wg} is 0.0728 N/m and σ_{hg} is 0.0184 N/m respectively. Use these data to estimate the interfacial tension between hexane and water. Compare this value to experimentally determined value of $\sigma_{wh} = 0.0511$ N/m.

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$$\sigma_{ab} = \sigma_{bg} + \sigma_{ag} - 2(\sigma_{ag}\sigma_{bg})^{\frac{1}{2}}$$

$$\sigma_{wh} = 0.0180 \text{ N/m}$$

This is 65% accuracy or 35% of the actual value.

$$Sp_{hw} = \sigma_{wg} - \sigma_{hg} - \sigma_{wh}$$



$$Sp_{hw} = \sigma_{wg} - \sigma_{hg} - \sigma_{wh} = 0.0033 \text{ N/m}$$

- $Sp_{hw} > 0$, hexane would spontaneously spread over the surface of water
- However, close to zero and so tendency is weak.
- Hexane is unlikely to form lens-shaped droplets.
- Spread out into a film, not aggressively covers the entire liquid surface.

Spontaneous spreading of liquid helium over the walls of a Dewar flask

