

The background features a complex network of blue lines and arrows. Some lines are solid, while others are dashed. The arrows indicate a flow or direction, often following curved paths. The overall aesthetic is technical and modern, typical of an engineering or scientific presentation.

CE407 SEPARATIONS

Lecture 09

Instructor: David Courtemanche

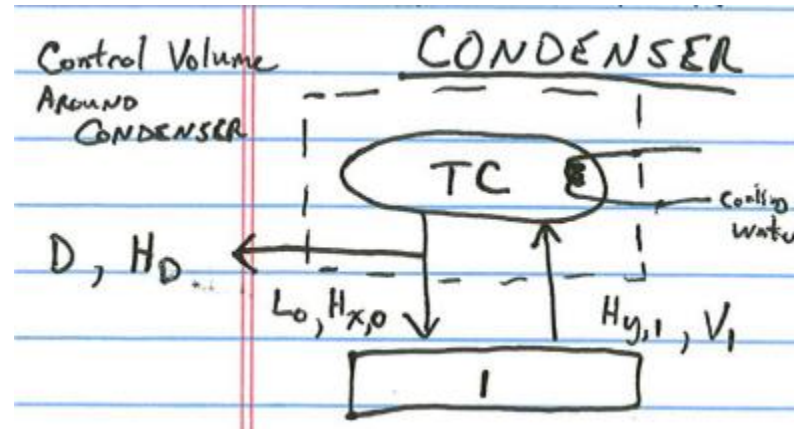
Continuous Distillation – Enthalpy Balances

McSH pp 694-701, pp 679, 682 eq 21.1 and 21.2

- How much energy does one need to remove via condenser and add via reboiler?

- **Condenser**

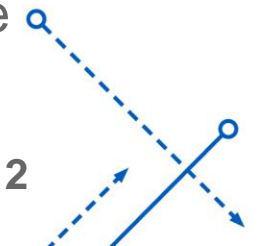
- Control Volume
around condenser



- Heat added = Enthalpy out – enthalpy in

$$\begin{aligned}
 -q_c &= DH_D + L_0H_{x,0} - V_1H_{y,1} \\
 &= (D + L_0)H_{x,0} - V_1H_{y,1}
 \end{aligned}$$

- Note: q_c is defined as the amount of heat **REMOVED** via the condenser and is a positive number. The enthalpy balance is the amount of heat added, therefore the term appears as $-q_c$ in the equation. The fact that $-q_c$ is therefore a negative term is consistent with the fact that we are removing heat to affect a phase change from vapor to liquid

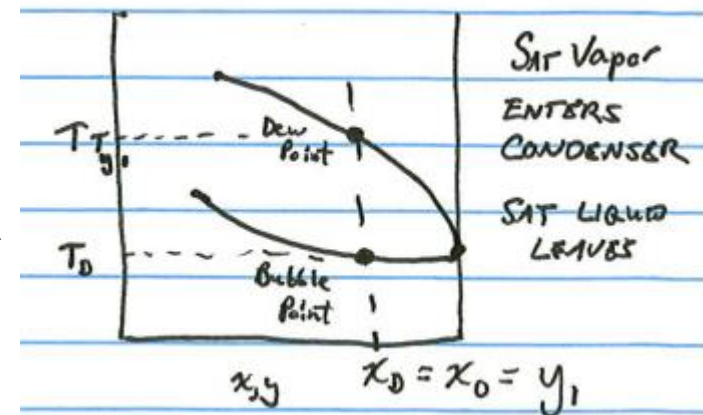
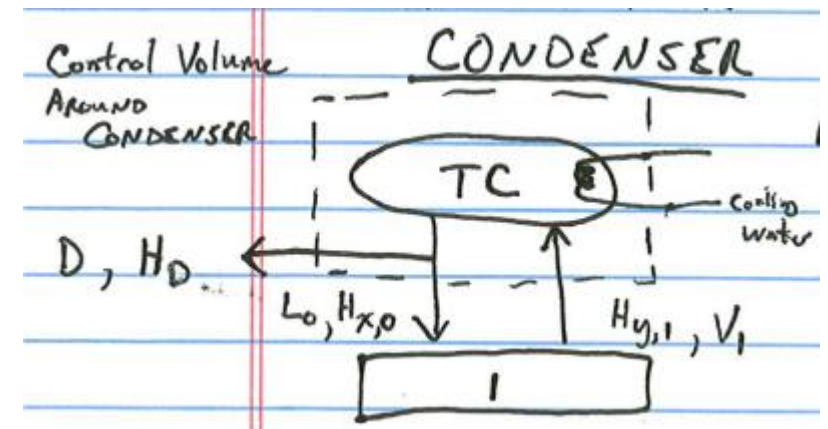


Enthalpy Balances: Condenser, continued

- $-q_c = (D + L_0)H_{x,0} - V_1H_{y,1}$
- But from mass balance we know that $V_1 = D + L_0$
- $-q_c = (D + L_0)H_{x,0} - (D + L_0)H_{y,1}$
- And from definition of reflux ratio $L_0 = D R$

$$-q_c = D(1 + R)(H_{x,0} - H_{y,1})$$

- It's not as simple as just looking at Heats of Vaporization
- You have the temperature changing from Dew Point to Bubble Point
- Heats of Vaporization are defined for a pure material at a given temperature (usually the normal boiling temperature for pure material) and we are condensing over a range of temperatures



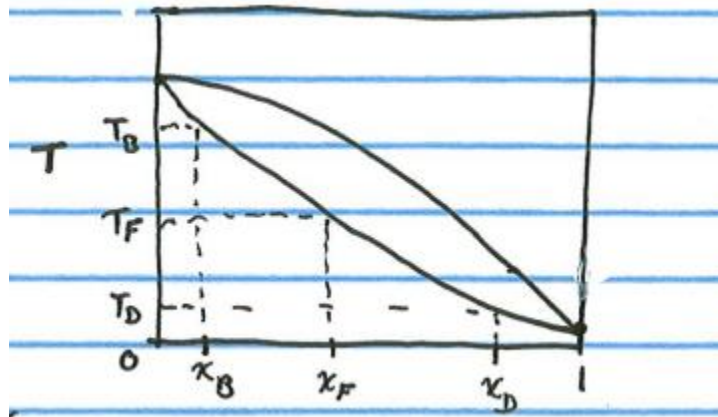
Enthalpy Balances: Reboiler

- Set up Control Volume around entire column

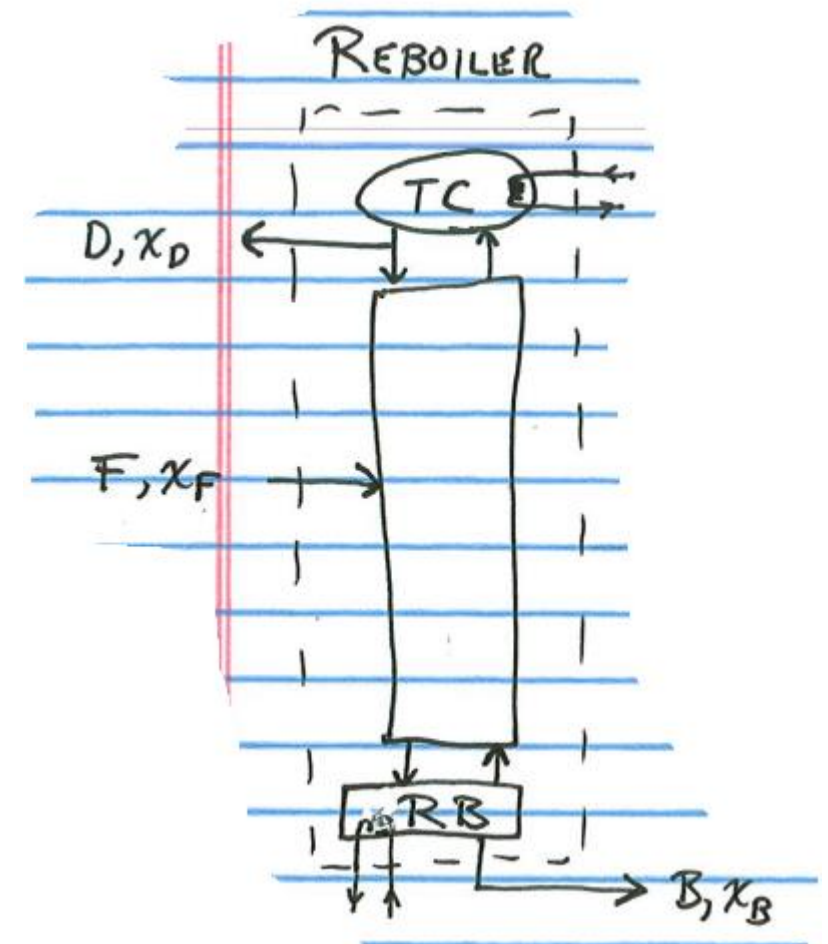
Heat added = Enthalpy out – enthalpy in

$$q_r - q_c = DH_D + BH_B - FH_F$$

- q_r is defined as heat ADDED via the Reboiler and is a positive number
- We will need to determine the temperature of each stream in order to calculate enthalpies

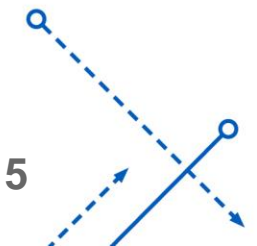


Txy diagram for saturated liquid feed case



Enthalpy Balances

- We have the temperature and composition of all streams
 - Calculate enthalpy of each stream
 - Start with enthalpy balance around condenser
 - Solve for q_c
 - Next evaluate enthalpy balance around entire column
 - Solve for q_r



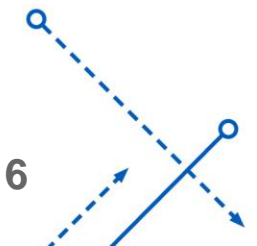
Enthalpy Balances – Heat Loads

Condenser

- Use q_c to determine rate of cooling water required for condenser

$$q_c = \dot{m}_{cw} C_{p H_2O}^L (T_{out} - T_{in})$$

- Where \dot{m}_{cw} is mass flow rate of cooling water
- $C_{p H_2O}^L$ is the heat capacity of liquid water
 - Typically $C_{p H_2O}^L \approx 1 \frac{cal}{g \text{ } ^\circ C}$
- T_{in} is the incoming temperature of the cooling water
- T_{out} is the outgoing temperature of the cooling water
 - this may be limited by thermal pollution concerns
 - of course, you don't want this stream to reach a boiling temperature



Enthalpy Balances – Heat Loads

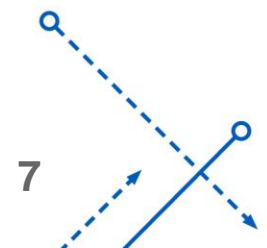
Reboiler

- Use q_r to determine rate of steam consumption required for reboiler

$$q_r = \dot{m}_{steam} \Delta H_{H_2O}^{vap}$$

- Where \dot{m}_{steam} is mass flow rate of steam
- $\Delta H_{H_2O}^{vap}$ is the heat of the steam phase change
- This equation assumes steam enters as saturated vapor and exits as saturated steam – consult steam tables for enthalpy change of steam that has varying quality values

	T (°C)	$P_{H_2O}^{sat}$ (kPa)	$\Delta H_{H_2O}^{vap}$ $\left(\frac{kJ}{kg}\right)$
	100	101.3	2257
~150 # steam	188	1200	1986

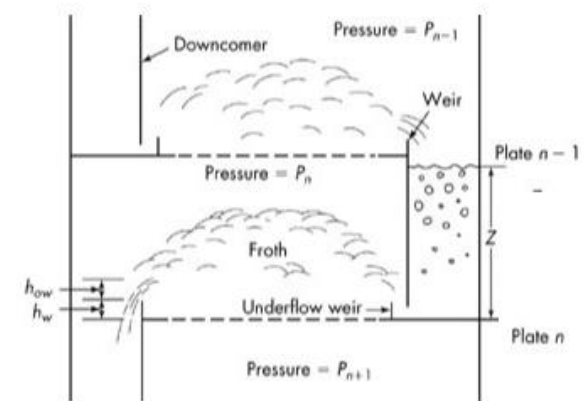
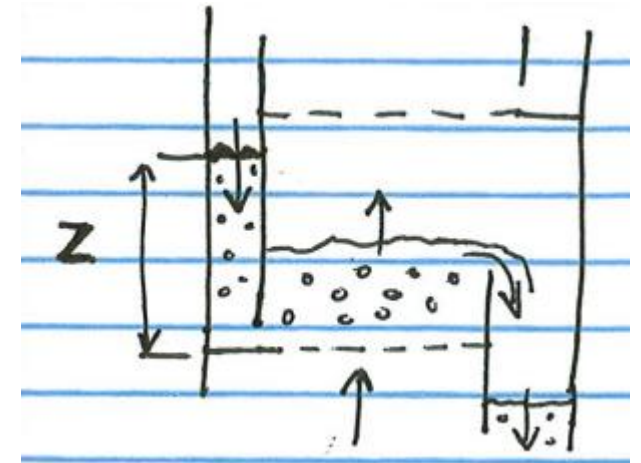


Continuous Distillation – Tower Design

McSH pp 701-712

Design of Sieve Plate Trays

- Pressure increases as you progress down the tower
- The pressure is needed to motivate the vapor through the holes in the tray and through the liquid held up on the tray
- Due to the ΔP , a column of liquid is held up in the downcomer (similar to a manometer) of height Z
- The text discusses methods for calculating Z , but we won't touch on it at this point
- The important point is that if Z exceeds the distance to the weir on the plate immediately above then the column experiences **FLOODING** and the plates won't function properly
- The diameter of the column must be specified such that the vapor velocity does not exceed the “**Flooding Velocity**”

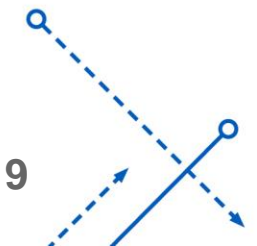
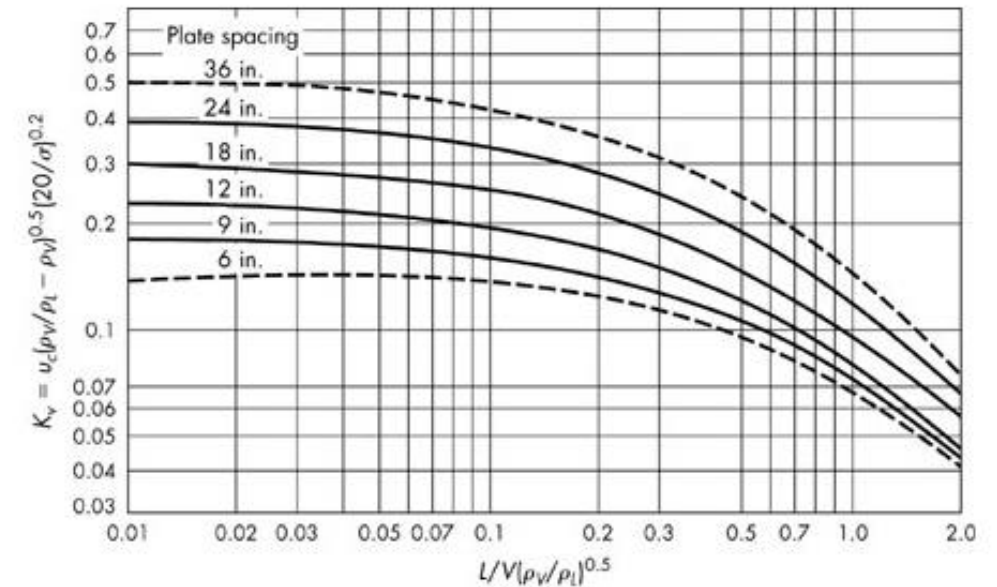


Flooding

- Figure 21.26 in McSH: Note that this is a log-log graph!
- Choose curve corresponding to your tray spacing
- Calculate $\frac{L_{mass}}{V_{mass}} \sqrt{\rho_V / \rho_L}$
 - Note that **L** and **V** are MASS flow rates
 - Note that ρ_V and ρ_L are MASS densities
 - **L/V** we have used in the past were molar flows
- Find K_V from chart
- Use equation 21.68 to calculate flooding velocity

$$u_c = u_{flooding} = K_V \sqrt{\frac{\rho_L - \rho_V}{\rho_V}} \left(\frac{\sigma}{20}\right)^{0.2}$$

- σ is the surface tension in dyn/cm, **u** is in ft/s on this graph
- A given chart will correspond to **u** in ft/s or m/s: *be sure you know which it is!*



Flooding

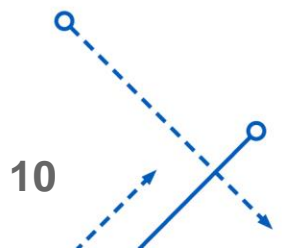
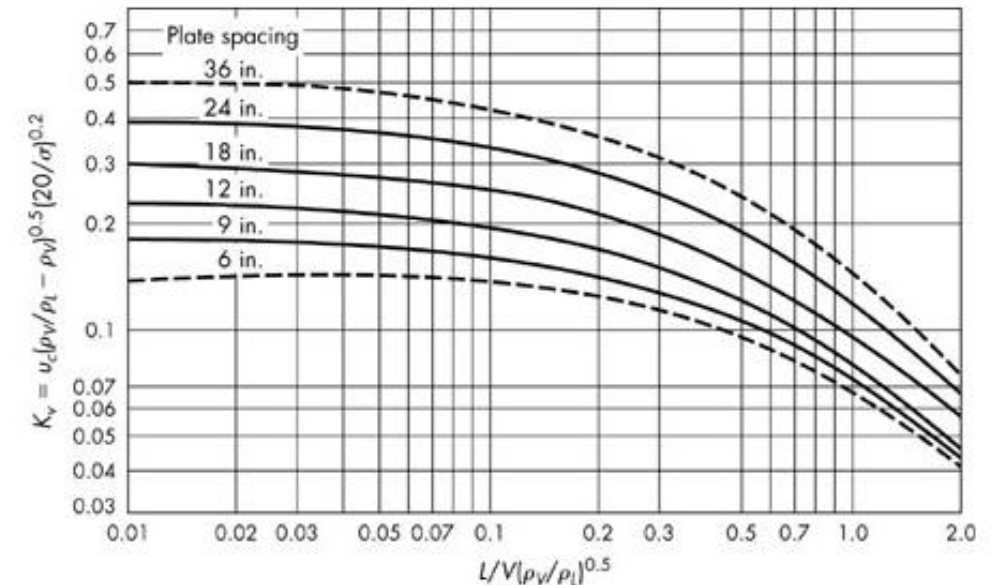
- Calculate $\frac{L_{mass}}{V_{mass}} \sqrt{\rho_V / \rho_L}$
 - Note that **L** and **V** are MASS flow rates
 - Note that ρ_V and ρ_L are MASS flow densities
 - **L/V** we have used in the past were molar flows

- $$\frac{L_{mass}}{V_{mass}} = \frac{L \overline{MW}_L}{V \overline{MW}_V}$$

- When we are evaluating at the top of the tower the flows are often close to pure light component and $\overline{MW}_L \approx \overline{MW}_V$

- Therefore $\frac{L_{mass}}{V_{mass}} \approx \frac{L}{V}$

- $\frac{L}{V} = \frac{R}{R+1}$ can be used for $\frac{L_{mass}}{V_{mass}}$ as long as we are aware that it only applies when $\overline{MW}_L \approx \overline{MW}_V$



Flooding, Continued

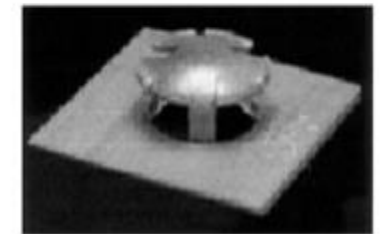
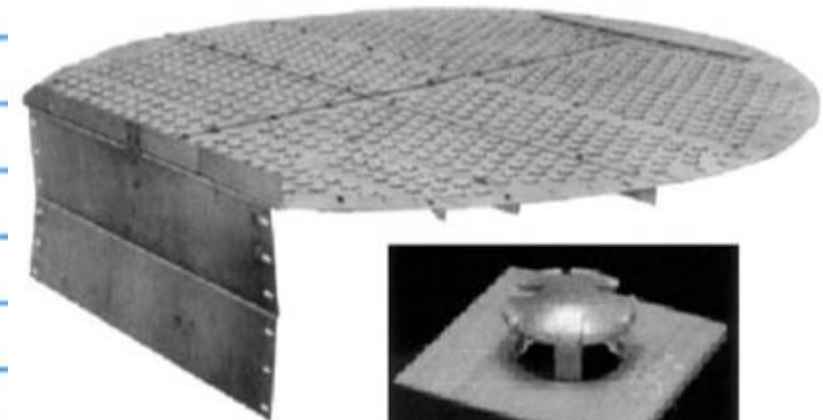
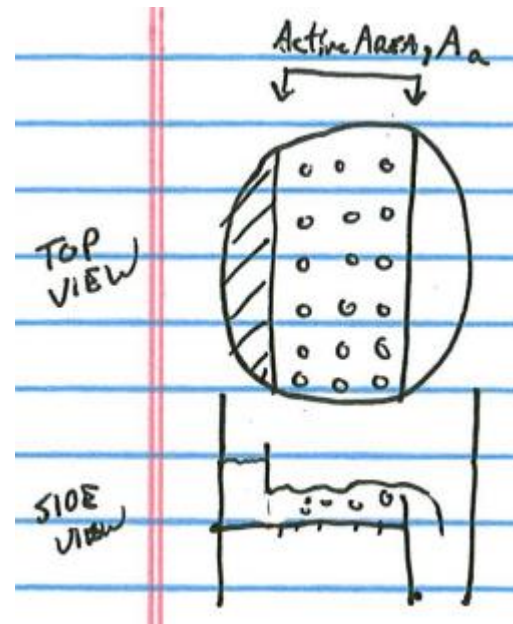
- Now we have a value for the a vapor velocity that will cause flooding.
- We will add a safety factor because we don't want to operate on the edge of flooding conditions
 - Various sources use different factors, let's choose $u \approx 0.7u_{flood}$

- Most of the plate is covered by holes

- $A = \text{Total Area} = \frac{\pi D^2}{4}$

- Downcomer, etc cover 15% of total area

- Net area for flow = **0.85A**

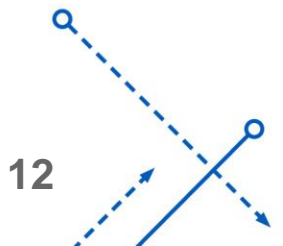


Flooding, Continued

$$V = u A_n \frac{P}{RT}$$

Molar vapor flow = (volumetric flow) * (moles/volume)

- From ideal gas law $PV = nRT$ you get $\frac{P}{RT} = \frac{n}{V} = \frac{\text{moles}}{\text{Volume}}$
- In this context V is just a volume
- $A_n = 0.85 \frac{\pi D^2}{4}$ net area for flow
- Steps
 - Calculate u_{flood} and apply safety factor
 - Use molar flow rate V to calculate A_n
 - Calculate required column diameter
- **Note: Column is most susceptible to flooding at the top of the tower. Use conditions corresponding to the top of the tower in this evaluation!**



Flooding Example

- We have a distillate flow rate of 3259 mol/minute of CCL_4 (light) and C_8H_{18} (heavy)
 - $x_D = 0.859$
- Reflux ratio = 2.55, Tray Spacing = 24"
- Flooding velocity is calculated at TOP of tower
- Problem Statement indicates that we should assume densities of the liquid and vapor to be that of pure CCL_4
 - Liquid density: $\rho_L = 1490 \frac{\text{kg}}{\text{m}^3}$
- Vapor density: By the Ideal Gas Law we can calculate molar volume

$$V = \frac{RT}{P} = \frac{83.14 \frac{\text{bar cm}^3}{\text{mol K}} (273.15 + 76.75) \text{ K}}{1.013 \text{ bar}} = 2.872 * 10^4 \frac{\text{cm}^3}{\text{mol}}$$

- Then the mass density can be calculated as

$$\rho_V = \frac{1 \text{ mol}}{2.872 * 10^4 \text{ cm}^3} * \frac{153.823 \text{ g}}{\text{mol}} * \frac{\text{kg}}{1000 \text{ g}} * \frac{10^6 \text{ cm}^3}{\text{m}^3} = 5.34 \frac{\text{kg}}{\text{m}^3}$$

- MW $\text{CCl}_4 = 153.823$



Flooding Example

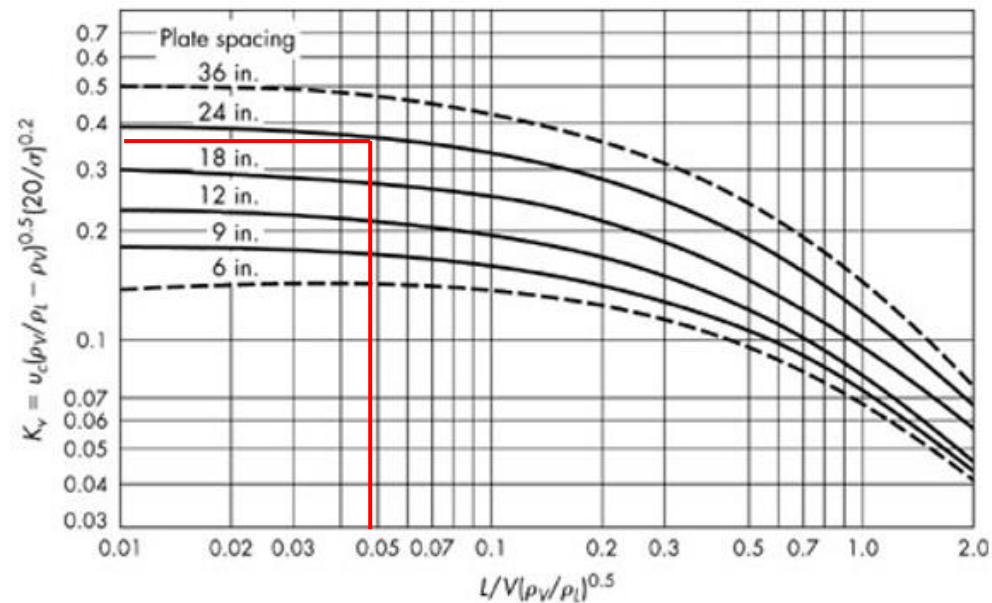
- Near the top of the tower (where the mole fractions are near 1.0), the average molecular weight of the liquid and the vapor are approximately equal at any location
- At the top of the tower $\frac{L_{mass}}{V_{mass}} = \frac{L * MW_L}{V * MW_V} \approx \frac{L}{V} = \frac{R}{R+1} = \frac{2.55}{2.55+1} = \mathbf{0.718}$

- $\frac{L_{mass}}{V_{mass}} \sqrt{\frac{\rho_V}{\rho_L}} = \mathbf{0.718} \sqrt{\frac{5.34 \frac{kg}{m^3}}{1490 \frac{kg}{m^3}}} = \mathbf{0.043}$

- $K_V = u_c \sqrt{\frac{\rho_V}{\rho_L - \rho_V}} \left(\frac{20}{\sigma}\right)^{0.2} = 0.37$

- Unless your surface tension is very different from water, the surface tension factor can be ignored

- $u_c = u_{flooding} = K_V \sqrt{\frac{\rho_L - \rho_V}{\rho_V}} = 0.37 \sqrt{\frac{1490 - 5.34}{5.34}} = 6.17 \frac{ft}{s}$



Flooding Example

- Vapor volumetric flow rate:
- V molar flow rate $V = L + D = (R + 1)D = 3.55 * 3259 \frac{\text{mol}}{\text{min}} = 11,569 \frac{\text{mol}}{\text{min}}$
- Molar density $\rho_V = \frac{P}{RT} = \frac{1.013 \text{ bar}}{83.14 \frac{\text{bar cm}^3}{\text{mol K}} (273.15 + 76.75) \text{ K}} = 3.482 * 10^{-5} \frac{\text{mol}}{\text{cm}^3} * \frac{10^6 \text{ cm}^3}{\text{m}^3} = 34.82 \frac{\text{mol}}{\text{m}^3}$
- $Q = \frac{V}{\rho_V} = \frac{11,569 \frac{\text{mol}}{\text{min}}}{34.82 \frac{\text{mol}}{\text{m}^3}} = 332.2 \frac{\text{m}^3}{\text{min}} * \frac{\text{min}}{60 \text{ s}} = 5.54 \frac{\text{m}^3}{\text{s}} * \left(\frac{\text{ft}}{0.305 \text{ m}} \right)^3 = 195.3 \frac{\text{ft}^3}{\text{s}}$

Flooding Example

- **Column diameter:**
- $Q = A_n * u = 0.85 \frac{\pi D^2}{4} * u$
 - (Volumetric flow rate = Area * velocity)
- Column is to operate at 50% of flooding velocity:
- $u = 0.5 * u_{flooding} = 0.5 * 6.17 \frac{ft}{s} = 3.09 \frac{ft}{s}$
- $Q = 195.3 \frac{ft^3}{s} = 0.85 \frac{\pi D^2}{4} * 3.09 \frac{ft}{s} = 2.06 \frac{ft}{s} * D^2$
- $D^2 = \frac{195.3 \frac{ft^3}{s}}{2.06 \frac{ft}{s}} = 94.67 ft^2$
- **$D = 9.7 ft$**