New Problems and new solutions are listed as new immediately after the solution number. These new problems are: 2A8, 2A10 parts c-e, 2A11, 2A12, 2A13, 2A14, 2C4, 2D1-part g, 2D3, 2D6, 2D7, 2D11, 2D13, 2D14, 2D20, 2D22, 2D23, 2D31, 2D32, 2E3, 2F4, 2G2, 2G3, 2H1, 2H3, 2H4, 2H5 and 2H6.

2.A1. Feed to flash drum is a liquid at high pressure. At this pressure its enthalpy can be calculated as a liquid. \( h(T_{F,P_{high}}) = c_{PLQ} (T_F - T_{ref}) \). When pressure is dropped the mixture is above its bubble point and is a two-phase mixture (It “flashes”). In the flash mixture enthalpy is unchanged but temperature changes. Feed location cannot be found from \( T_F \) and \( z \) on the graph because equilibrium data is at a lower pressure on the graph used for this calculation.


2.A3. The liquid is superheated when the pressure drops, and the energy comes from the amount of superheat.


2.A6. In a flash drum separating a multicomponent mixture, raising the pressure will:
   i. Decrease the drum diameter and decrease the relative volatilities.  \( \text{Answer is i.} \)

   a. At 100°C and a pressure of 200 kPa what is the K value of n-hexane? 0.29
   b. As the pressure increases, the K value
      a. increases, b. decreases, c. stays constant \( \text{b} \)
   c. Within a homologous series such as light hydrocarbons as the molecular weight increases, the K value (at constant pressure and temperature)
      a. increases, b. decreases, c. stays constant \( \text{b} \)
   d. At what pressure does pure propane boil at a temperature of -30°C? 160 kPa
a. The answer is 3.5 to 3.6
b. The answer is 36°C
c. This part is new in 4th ed. 102°C

2.A10. Parts c, d, and e are new in 4th ed.  
a. 0.22;  b. No;  c. From y-x plot for Methanol x = 0.65, \( y_M = 0.85 \); thus, \( y_W = 0.15 \).  
d. \( K_M = 0.579/0.2 = 2.895 \), \( K_W = (1 - 0.579)/(1 - 0.2) = 0.52625 \).  
e. \( \alpha_{M,W} = K_M/K_W = 2.895/0.52625 = 5.501 \).

2.A11. New problem in 4th edition. Because of the presence of air this is not a binary system. Also, it is not at equilibrium.

2.A12. New problem in 4th edition. The entire system design includes extensive variables and intensive variables necessary to solve mass and energy balances. Gibbs phase rule refers only to the intensive variables needed to set equilibrium conditions.

2A13. New problem in 4th edition. Although V is an extensive variable, V/F is an intensive variable and thus satisfies Gibbs phase rule.

2A14. New problem in 4th edition. 1.0 kg/cm² = 0.980665 bar = 0.96784 atm.
Source: http://www.unit-conversion.info/pressure.html

2.B1. Must be sure you don’t violate Gibbs phase rule for intensive variables in equilibrium.
Examples:
- \( F, z, T_{drum}, P_{drum} \)  
- \( F, z, y, P_{drum} \)  
- \( F, z, x, P_{drum} \)  
- \( F, z, y, P_{drum} \)  
- \( F, z, x, T_{drum} \)  
- Drum dimensions, \( z, F_{drum}, P_{drum} \)  
- Drum dimensions, \( z, y, P_{drum} \)  
- etc.

2.B2. This is essentially the same problem (disguised) as problem 2-D1c and e but with an existing (larger) drum and a higher flow rate.

With \( y = 0.58, x = 0.20, \) and \( V/F = 0.25 \) which corresponds to 2-D1c.

If \( F = 1000 \frac{lb\ mole}{hr} \), \( D = .98 \) and \( L = 2.95 \) ft from Problem 2-D1e.

Since \( D \propto \sqrt{V} \) and for constant \( V/F, V \propto F \), we have \( D \propto \sqrt{F} \).

With \( F = 25,000 \):

\[
\sqrt{\frac{F_{new}}{F_{old}}} = 5, \quad D_{new} = 5 D_{old} = 4.90, \text{ and } L_{new} = 3 D_{new} = 14.7.
\]

Existing drum is too small.
Feed rate drum can handle: \( F \propto D^2 \). \( \frac{F_{\text{existing}}}{1000} = \left( \frac{D_{\text{existing}}}{.98} \right)^2 = \left( \frac{4}{.98} \right)^2 \) gives

\[ F_{\text{existing}} = 16,660 \text{ lbmol/h} \]

**Alternatives**

a) Do drums in parallel. Add a second drum which can handle remaining 8340 lbmol/h.
b) Bypass with liquid mixing

c) Look at Eq. (2-62), which becomes

\[ V = \sqrt{\frac{3K_{\text{drum}} 3600}{\rho_v}} \left( \rho_L - \rho_v \right) \rho_v \]

Bypass reduces \( V \)

c1) \( K_{\text{drum}} \) is already 0.35. Perhaps small improvements can be made with a better demister → Talk to the manufacturers.
c2) \( \rho_v \) can be increased by increasing pressure. Thus operate at higher pressure. Note this will change the equilibrium data and raise temperature. Thus a complete new calculation needs to be done.
d) Try bypass with vapor mixing.
e) Other alternatives are possible.

2.C2.

\[ \frac{V}{F} = \left[ \frac{-z_A}{(K_B - 1)} - \frac{z_B}{(K_A - 1)} \right] \]

2.C5. a. Start with \( x_i = \frac{Fz_i}{L + VK_i} \) and let \( V = F - L \)

\[ x_i = \frac{Fz_i}{L + (F - L)K_i} \text{ or } x_i = \frac{L}{F} + \left( 1 - \frac{L}{F} \right) K_i \]

\( \rho_v \)
Then \( y_i = K_i x_i = \frac{K_i z_i}{L + \left( \frac{L}{F} \right) K_i} \)

From \( \sum y_i - \sum x_i = 0 \) we obtain \( \sum \frac{(K_i - 1) z_i}{L + \left( \frac{L}{F} \right) K_i} = 0 \)

2.C4. *New Problem.* Prove that the intersection of the operating and \( y = x \) lines for binary flash distillation occurs at the mole fraction of the feed.

**SOLUTION:**

\[
y = \frac{L}{V} y + \frac{F}{V} z
\]

Rearrange:

\[
y \left[ 1 + \frac{L}{V} \right] = \frac{F}{V} z \quad \text{or} \quad y \left[ \frac{V + L}{V} \right] = \frac{F}{V} z
\]

Since \( V + L = F \), the result is \( y = z \) and therefore \( x = y = z \) \((2-18)\)

The intersection is at the feed composition.

2.C7.

\[
\sum \frac{z_i}{1 + (K_i - 1) \frac{V}{F}} - 1 = f \left( \frac{V}{F} \right)
\]

From data in Example 2-2 obtain:

<table>
<thead>
<tr>
<th>( V/F )</th>
<th>0</th>
<th>.1</th>
<th>.2</th>
<th>.3</th>
<th>.4</th>
<th>.5</th>
<th>.6</th>
<th>.7</th>
<th>.8</th>
<th>.9</th>
<th>1.0</th>
</tr>
</thead>
<tbody>
<tr>
<td>( f )</td>
<td>0</td>
<td>-.09</td>
<td>-.1</td>
<td>-.09</td>
<td>-.06</td>
<td>-.007</td>
<td>.07</td>
<td>.16</td>
<td>.3</td>
<td>.49</td>
<td>.77</td>
</tr>
</tbody>
</table>

![Graph](image_url)
2.C8. Derivation of Eqs. (2-62) and (2-63). Overall and component mass balances are,

\[ F = V + L_1 + L_2 \text{ and } Fz_i = L_1 x_{i,1} + L_2 + x_{i,1} + V y_i \]

Substituting in Eqs. (2-60b) and 2-60c)

\[ Fz_i = L_1 K_{i,1} x_{i,1} + L_2 x_{i,1} + VK_{i,1V} x_{i,1} \]

Solving,

\[ x_{i,1} = \frac{Fz_i}{L_1 K_{i,1} + L_2 + VK_{i,1V}} \]

Dividing numerator and denominator by \( F \) and collecting terms.

\[ x_{i,liq2} = \frac{z_i}{1 + \left(K_{i,1} - 1\right) \frac{L_1}{F} + \left(K_{i,1V} - 1\right) \frac{V}{F}} \]

Since \( y_i = K_{i,1V} x_{i,1} \), \( y_i = \frac{K_{i,1V} z_i}{1 + \left(K_{i,1} - 1\right) \frac{L_1}{F} + \left(K_{i,1V} - 1\right) \frac{V}{F}} \]

Stoichiometric equations,

\[ \sum_{i=1}^{C} x_{i,1} = 1, \sum_{i=1}^{C} y_i = 1, \text{ thus, } \sum_{i=1}^{C} y_i - \sum_{i=1}^{C} x_{i,1} = 0 \]

which becomes

\[ \sum_{i=1}^{C} \left(\frac{K_{i,1} - 1}{1 + \left(K_{i,1} - 1\right) \frac{L_1}{F} + \left(K_{i,1V} - 1\right) \frac{V}{F}} \right) z_i = 0 \] (2-62)

Since \( x_{i,liq} = K_{i,1} x_{i,1} \), we have

\[ x_{i,liq1} = \frac{K_{i,1} z_i}{1 + \left(K_{i,1} - 1\right) \frac{L_1}{F} + \left(K_{i,1V} - 1\right) \frac{V}{F}} \]

In addition,

\[ \sum_{i=1}^{C} x_{i,liq1} - \sum_{i=1}^{C} x_{i,liq2} = 0 = \sum_{i=1}^{C} \left(\frac{K_{i,1} - 1}{1 + \left(K_{i,1} - 1\right) \frac{L_1}{F} + \left(K_{i,1V} - 1\right) \frac{V}{F}} \right) z_i \] (2-63)

2.D1. a. \( V = (0.4)100 = 40 \) and \( L = F - V = 60 \) kmol/h

Slope op. line = \( -L/V = -3/2, y = x = z = 0.6 \)

See graph. \( y = 0.77 \) and \( x = 0.48 \)
b. \( V = (0.4)(1500) = 600 \) and \( L = 900 \). Rest same as part a.
c. Plot \( x = 0.2 \) on equil. Diagram and \( y = x = z = 0.3 \). \( y_{\text{intercept}} = zF/V = 1.2 \)

\( V/F = z/1.2 = 0.25 \). From equil \( y = 0.58 \).
d. Plot \( x = 0.45 \) on equilibrium curve.

Slope = \( -\frac{L}{V} = -\frac{F - V}{V} = -\frac{1 - V/F}{V/F} = -\frac{0.8}{0.2} = -4 \)
Plot operating line, \( y = x = z \) at \( z = 0.51 \). From mass balance \( F = 37.5 \ \text{kmol/h} \).

e. Find Liquid Density.

\[
\overline{\text{MW}}_L = x_m (\text{MW}_m) + x_w (\text{MW}_w) = (0.2)(32.04) + (0.8)(18.01) = 20.82
\]

Then,

\[
\overline{\text{V}}_L = x_m \frac{\text{MW}_m}{\rho_m} + x_w \frac{\text{MW}_w}{\rho_w} = 0.2 \left( \frac{32.04}{0.7914} \right) + 0.8 \left( \frac{18.01}{1.00} \right) = 22.51 \ \text{ml/mol}
\]

\[
\rho_L = \frac{\overline{\text{MW}}_L}{\overline{\text{V}}_L} = \frac{20.82}{22.51} = 0.925 \ \text{g/ml}
\]

Vapor Density: \( \rho_v = \frac{p(\text{MW})_{v,avg}}{RT} \) (Need temperature of the drum)

\[
\overline{\text{MW}}_v = y_m (\text{MW})_m + y_w (\text{MW})_w = 0.58(32.04) + 0.42(18.01) = 26.15 \ \text{g/mol}
\]

Find Temperature of the Drum \( T \): From Table 3-3 find \( T \) when \( y = 0.58, x = 20, T = 81.7^\circ C = 354.7 \text{K} \)

\[
\rho_v = (1 \text{ atm})(26.15 \text{ g/mol}) \left[ \frac{82.0575 \text{ ml atm}}{\text{ mol K}} \right] = 8.98 \times 10^{-4} \text{ g/ml}
\]

Find Permissible velocity:

\[
u_{per} = K_{d,m} \sqrt{(\rho_L - \rho_v)/\rho_v}, \quad K_{d,m} = \exp \left[ A + B(\epsilon n_F)_m + C(\epsilon n_F)_w + D(\epsilon n_F) + E(\epsilon n_F) \right]
\]

\[
V = \left( \frac{V}{F} \right) F = 0.25 \times 1000 = 250 \ \text{lbmol/h}, \quad W_v = V \left( \overline{\text{MW}}_v \right) = 250 \left( 26.15 \frac{\text{lb}}{\text{lbmol}} \right) = 6537.5 \ \text{lb/h}
\]

\[
L = F - V = 1000 - 250 = 750 \ \text{lbmol/h}, \quad \text{and} \quad W_L = (L) \left( \overline{\text{MW}}_L \right) = (750)(20.82) = 15,615 \ \text{lb/h}
\]
\[ F_v = \frac{W_L}{W_v} \sqrt{\frac{\rho_v}{\rho_L}} = \left( \frac{15615}{6537.5} \right) \sqrt{\frac{8.89 \times 10^{-4}}{0.925}} = 0.0744, \text{ and } \ell n(F_h) = -2.598 \]

Then \( K_{\text{drum}} = 0.442 \), and \( u_{\text{perm}} = 0.442 \sqrt{\frac{925-8.98 \times 10^{-4}}{8.98 \times 10^{-4}}} = 14.19 \text{ ft/s} \)

\[ A_{\text{cs}} = \frac{\sqrt{\text{MW}_v}}{u_{\text{perm}} 3600 \rho_v} = \frac{250 (26.15) (454 \text{ g/lb})}{(14.19) (3600) (8.98 \times 10^{-4} \text{ g/ml}) (28316.85 \text{ ml/ft}^3)} = 2.28 \text{ ft}^2. \]

\[ D = \sqrt{4A_{\text{cs}}/\pi} = 1.705 \text{ ft}. \text{ Use 2 ft diameter. L ranges from } 3 \times D = 6 \text{ ft to } 5 \times D = 10 \text{ ft} \]

Note that this design is conservative if a demister is used.

f. Plot \( T \) vs \( x \) from Table 3-3. When \( T = 77 \degree C, x = 0.34, y = 0.69 \). This problem is now very similar to 3-D1c. Can calculate \( V/F \) from mass balance, \( F_z = L_x + V_y \). This is

\[ F_z = (F-V)x + Vy \text{ or } \frac{V}{F} = \frac{z-y}{y-x} = \frac{0.4-0.34}{0.69-0.34} = 0.17 \]

g. Part g is a new problem. \( V = 16.18 \text{ mol/h, L = 33.82, y= 0.892, x = 0.756}. \)

2-D2. Work backwards. Starting with \( x_2 \), find \( y_2 = 0.62 \) from equilibrium. From equilibrium point plot op. line of slope \( = -(L/V)_2 = -\left(1 - \frac{V}{F}\right)_2 = -3/7 \). Find \( z_2 = 0.51 = x_1 \) (see Figure). From equilibrium, \( y_1 = 0.78 \). For stage 1, \( \frac{V}{F} = \frac{z_1-x_1}{y_1-x_1} = \frac{0.55-0.51}{0.78-0.51} = 0.148 \).

<table>
<thead>
<tr>
<th>x ethane</th>
<th>y ethane</th>
<th>T °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>0</td>
<td>63.19</td>
</tr>
<tr>
<td>.025</td>
<td>.01610</td>
<td>56.18</td>
</tr>
<tr>
<td>.05</td>
<td>.2970</td>
<td>49.57</td>
</tr>
<tr>
<td>.10</td>
<td>.5060</td>
<td>37.57</td>
</tr>
<tr>
<td>.15</td>
<td>.6503</td>
<td>27.17</td>
</tr>
<tr>
<td>.20</td>
<td>.7492</td>
<td>18.26</td>
</tr>
<tr>
<td>.25</td>
<td>.8175</td>
<td>10.64</td>
</tr>
<tr>
<td>.30</td>
<td>.8652</td>
<td>4.11</td>
</tr>
<tr>
<td>1.0</td>
<td>1.0</td>
<td>-37.47</td>
</tr>
</tbody>
</table>

b. See Figure. a. If 1 bubble of vapor product (V/F = 0) vapor product, vapor y_E = 0.7492 (highest) liquid x_E = z_E = 0.20 (highest) and T = 18.26 °C. If 1 drop of liquid product (V/F = 1) y_E = z_E = 0.20 (lowest), x_E = 0.035, T (by linear interpolation) ~ 56.18 + [(49.57 – 56.18)/(.297 -.161)][.2 – 0.16] = 54.2 °C (highest).

c. See figure. Slope = -L/V = - (1 – V/F)/(V/F) = -.6/.4 = -1.5. x_E = 0.12, y_E = 0.57, T = 33.4°C.

d. From equilibrium data y_E = 0.7492. For an F = 1, L = 1 – V, Ethane balance: .2L = 1(.3) – 0.7492 V. Solve 2 equations: V/F = 0.1821. Can also find V/F from slope of operating line.

e. If do linear interpolation on equilibrium data, x = 0.05 + (.45-49.57)(.05 -0.05)/(37.57 – 49.57) = 0.069. From equilibrium plot y = 0.375.

Mass balance for basis F = 1, L = 1 – V and 0.069 L = 0.18 – 0.375 V. Solve simultaneously, V/F = 0.363.

2.D4. **New problem in 3rd edition.** Highest temperature is dew point (V/F = 0)

Set \( z_i = y_i \).

\[ K_i = y_i / x_i \]

Want \( \sum x_i = \sum y_i / K_i = 1.0 \)

\[ K_{ref \ (T新技术)} = K_{ref \ (T旧)} \left( \sum \left( y_i / K_i \right) \right) \]

If pick C4 as reference: First guess \( K_{butane} = 1.0 \), \( T = 41 \text{°C} \): \( K_{C3} = 3.1, K_{C6} = 0.125 \)

\[ \sum \frac{y_i}{K_i} = \frac{.2}{3.1} + \frac{.35}{1.0} + \frac{.45}{.125} = 4.0145 \quad \text{T too low} \]

Guess for reference: \( K_{C4} = 4.014 \), \( T = 118 \text{°C} \): \( K_{C3} = 8.8, K_{C6} = .9 \)

\[ \sum \frac{y_i}{K_i} = \frac{.2}{8.8} + \frac{.35}{4.0145} + \frac{.45}{.9} = 0.6099 \]

\[ K_{C4,NEW} = 4.0145(.6099) = 2.45, T = 85 \text{°C} : K_{C2} = 6.0, K_{C6} = 0.44 \]
\[ \sum \frac{y_i}{K_i} = \frac{2}{6} + \frac{.35}{2.45} + \frac{.45}{.44} = 1.20 \]

\[ K_{C4,NEW} = 2.45 \times 1.2 = 2.94, \; T = 96^\circ C: \; K_{C3} = 6.9, \; K_{C6} = 0.56 \]

\[ \sum \frac{y_i}{K_i} = \frac{2}{6.9} + \frac{.35}{2.94} + \frac{.45}{.56} = 0.804 \Rightarrow \text{Gives } 84^\circ C \]

Use 90.5° → Avg last two T \( K_{C4} = 2.7, \; K_{C3} = 6.5, \; K_{C6} = 0.49 \)

\[ \sum \left( \frac{y_i}{K_i} \right) = \frac{2}{6.5} + \frac{.35}{2.7} + \frac{.45}{0.49} = 1.079, \; T \sim 87-88^\circ C \]

Note: Hexane probably better choice as reference.

2.D5. a)

\begin{align*}
V_1 & = F_2 \\
F_1 & = 1000 \\
x_1 & = 0.30 \\
z_1 & = 0.55 \\
& \\
\end{align*}

\[ \text{Plot 1st Op line.} \]

\[ y = x = z = 0.55 \text{ to } x_1 = 0.3 \text{ on eq. curve (see graph)} \]

\[ \text{Slope } = -\frac{L}{V} = \frac{0.55 - 0.80}{0.55} = -0.454545 \]

\[ L_1 + V_1 = F_1 = 1000 \]

\[ V_1 = 687.5 \text{ kmol/h} = F_2 \]

\[ \frac{V}{F} = \frac{687.5}{1000} = 0.6875 \]

b) \( y_i = -\frac{L}{V} x_i + \frac{F}{V} z \) Plot 1st Op line.

\[ y_1 = 0.66 = z_2 \]

\[ y = x = z = 0.55 \text{ to } x_1 = 0.3 \text{ on eq. curve (see graph)} \]

\[ \text{Slope } = -\frac{L}{V} = \frac{0.55 - 0.80}{0.55} = -0.454545 \]

\[ L_1 + V_1 = F_1 = 1000 \]

\[ V_1 = 687.5 \text{ kmol/h} = F_2 \]

\[ \frac{V}{F} = \frac{687.5}{1000} = 0.6875 \]

c) Stage 2 \( \frac{V}{F} = 0.25 \), \( \frac{L}{V} = -\frac{0.75F}{0.25F} = -3 \), \( y = x = z = 0.66 \). Plot Op line

\[ \text{At } x = 0, \; y = \frac{z}{(V/F)} = \frac{0.66}{0.25} = 2.64. \; \text{At } y = 0, \; x_2 = \frac{F}{L} = \frac{z}{L/F} = \frac{0.66}{0.75} = 0.88 \]

\[ \text{From graph } y_2 = 0.82, \; x_2 = 0.63 \]

\[ V_2 = \frac{V}{F} F_2 = (0.25)687.5 = 171.875 \text{ kmol/h} \]
2.D6. New problem in 4th ed. a.) The answer is $VP = 19.30$ mm Hg

$$\log_{10}(VP) = 6.8379 - \frac{1310.62}{100 + 136.05} = 1.2856$$

b.) The answer is $K = 0.01693$. $K = \frac{VP}{P_{tot}} = \frac{19.30}{1.5(760)}$

Part a. Drum 1: $V_1/F_1 = 0.3$, Slope op line $= -L/V = -7/3 = -2.33$, $y=x=z_1 = 0.46$. $L_1 = F_2 = 70$.
From graph $x_1 = z_2 = 0.395$
Drum 2: $V_1/F_1 = 30/70$, Slope op line $= -L/V = -7/3$, $y=x=z_2 = 0.395$. $L_1 = F_2 - V_2 = 40$.
From graph $x_2 = 0.263$

Part b. Single drum: $V/F = 0.6$, Slope op line $= -L/V = -40/60 = -2/3$, From graph $x = 0.295$.
More separation with 2 drums.
2.D8. Use Rachford-Rice eqn: 
\[ f\left(\frac{V}{F}\right) = \sum \frac{(K_i - 1)z_i}{1 + (K_i - 1)\frac{V}{F}} = 0 \] . Note that 2 atm = 203 kPa.

Find \( K_i \) from DePriester Chart: \( K_1 = 73, \ K_2 = 4.1 \ K_3 = .115 \)

Converge on \( V/F = .076, \ V = F(V/F) = 152 \) kmol/h, \( L = F - V = 1848 \) kmol/h.

From \( x_i = \frac{z_i}{1 + \frac{V}{F}(K_i - 1)} \) we obtain \( x_1 = .0077, \ x_2 = .0809, \ x_3 = .9113 \)

From \( y_i = K_i x_i \), we obtain \( y_1 = .5621, \ y_2 = .3649, \ y_3 = .1048 \)

2.D9. Need \( h_f \) to plot on diagram. Since pressure is high, feed remains a liquid

\[ h_f = \overline{C_p} (T_f - T_{ref}) \, \text{from chart} \]

\[ \overline{C_p} = C_{p_{\text{EtOH}}} x_{\text{EtOH}} + C_{p_{\text{water}}} x_{\text{water}} \]

Where \( x_{\text{EtOH}} \) and \( x_{\text{water}} \) are mole fractions. Convert weight to mole fractions.

Basis: 100 kg mixture: 30 kg EtOH = \( \frac{30}{46.07} = 0.651 \) kmol

70 kg water = 70/18.016 = 3.885 Total = 4.536 kmol
Avg. MW = \frac{100}{4.536} = 22.046 \quad \text{Mole fracs:} \quad x_E = \frac{0.6512}{4.536} = 0.1435, \quad x_w = 0.8565.

Use \( C_{p,\text{avg}} \) at 100°C as an average \( C_p \) value.

\[
\overline{C_{p_L}} = 37.96 (0.1435) + 18.0 (0.8565) = 20.86 \quad \text{kcal/kmol °C}
\]

Per kg this is \( \frac{\overline{C_{p_L}}}{\text{MW}_{\text{avg}}} = \frac{20.86}{22.046} = 0.946 \quad \text{kcal/kg °C} \)

\( h_f = 0.946 (2000) = 189.2 \quad \text{kcal/kg} \)

which can now be plotted on the enthalpy composition diagram.

Obtain \( T_{\text{drum}} \approx 88.2°C, \quad x_E = 0.146, \quad \text{and} \quad y_E = 0.617. \)

For \( F = 1000 \) find \( L \) and \( V \) from \( F = L + V \) and \( Fz = Lx + Vy \)

which gives \( V = 326.9, \) and \( L = 673.1 \)

Note: If use wt. fracs. \( \overline{C_{p_L}} = 23.99 \) & \( \frac{\overline{C_{p_L}}}{\text{MW}_{\text{avg}}} = 1.088 \) and \( h_f = 217.6. \) All wrong.

2.D.10 Solution 400 kPa, 70°C

\( z_{c_4} = 35 \quad \text{Mole % n-butane} \quad x_{c_6} = 0.7 \)

From DePriester chart \( K_{c_3} = 5, \quad K_{c_4} = 1.9, \quad K_{c_6} = 0.3 \)

Know \( y_i = K_i x_i, \quad x_i = \frac{z_i}{1 + (K_i - 1) \frac{V}{F}}, \quad \sum x_i = \sum y_i = 1 = \sum z_i \)
R.R. \[ \sum \frac{(K_i - 1)z_i}{1 + (K_i - 1)} \frac{V}{F} = 0 \]
\[ z_{c3} = 1 - z_{c6} - z_{c4} = 0.65 - z_{c6} \]

C6: \[ 0.7 = \frac{z_{c6}}{1 + (K_{c6} - 1)} \frac{V}{F} = \frac{z_{c6}}{1 - 0.7} \frac{V}{F} \implies z_{c6} = 0.7 \left( 1 - 0.7 \frac{V}{F} \right), \quad z_{c6} = 0.7 - 0.49 \frac{V}{F} \]

RR Eq:
\[ \frac{4(0.65 - z_{c6})}{1 + 4 \frac{V}{F}} + \frac{0.9(0.35)}{1 + 0.9 \frac{V}{F}} - \frac{0.7z_{c6}}{1 - 0.7 \frac{V}{F}} = 0 \]

2 equations & 2 unknowns. Substitute in for \( z_{c6} \). Do in Spreadsheet. Use Goal – Seek to find \( V/F \). \( V/F = 0.594 \) when R.R. equation \( 0.000881 \).

\[ z_{c6} = 0.7 - 0.49 \frac{V}{F} = 0.7 - (0.49)(0.594) = 0.40894 \]


\[ K_E = \frac{y_E}{x_E} \quad \text{and} \quad K_P = \frac{y_P}{x_P} \]
Since \( y_p = 1 - y_E \) and \( x_p = 1 - x_E \), \( K_P = (1 - y_E)/(1 - x_E) \).

Thus, 2 eqs and 2 unknowns. Solve for \( y_E \) and \( x_E \).

\[ x_E = \frac{(1 - K_P)}{(K_E - K_P)} \quad \text{and} \quad y_E = K_E x_E = K_E \left( 1 - \frac{x_E}{K_E} \right) \]

\( x_E = (1 - 0.56) / (2.2 - 0.56) = 0.268 \) and \( y_E = K_E x_E = (2.2)(0.268) = 0.590 \)

Check: \( x_p = 1 - x_E = 1 - 0.268 = 0.732 \) and \( y_p = 1 - y_E = 1 - 0.590 = 0.410 \)

\( K_P = y_p/x_p = 0.410/0.732 = 0.56 \) OK

2.D12. For problem 2.D1c, plot \( x = 0.2 \) on equilibrium diagram with feed composition of 0.3.

The resulting operating line has a y intercept \( z/(V/F) = 1.2 \). Thus \( V/F = 0.25 \) (see figure in Solution to 2.D1) Vapor mole fraction is \( y = 0.58 \).

Find Liquid Density:
\[ \frac{\rho_L}{\rho_m} = x_m \left( \frac{MW_m}{\rho_m} \right) + x_w \left( \frac{MW_w}{\rho_w} \right) = \frac{0.2}{32.04} + \frac{0.8}{18.01} = 0.2251 \text{ ml/mol} \]

\[ \rho_L = \frac{\rho_L}{\rho_m} MW_L = 0.2251 \times 20.82 = 0.925 \text{ g/ml} \]

Find Vapor Density:
\[ \rho_v = \frac{p(\frac{MW}{\rho_m})}{RT} \]

\[ \frac{MW_v}{\rho_m} = y_m (\frac{MW_m}{\rho_m}) + y_w (\frac{MW_w}{\rho_w}) = 0.58(32.04) + 0.42(18.01) = 26.15 \text{ g/mol} \]

Find Temperature of the Drum T:

From Table 2-7 find \( T \) corresponding to \( y = 0.58 \), \( x = 20 \), \( T = 81.7^\circ \text{C} = 354.7 \text{ K} \)

\[ \rho_v = (1 \text{ atm})(26.15 \text{ g/mol})/\left[ \left[ 82.0575 \text{ g atm mol}^{-1} \text{ K} \right] (354.7 \text{ K}) \right] = 8.98 \times 10^{-4} \text{ g/ml} \]

Find Permissible Velocity:
\[ u_{perm} = K_{dum} \sqrt{\frac{(\rho_L - \rho_v)}{\rho_v}} \]
\[ K_{\text{drum, horizontal}} = 1.25 \times K_{\text{drum, vertical}} = \left\{ \exp \left[ A + B \left( \ell n F_v \right) + C \left( \ell n F_v \right)^2 + D \left( \ell n F_v \right)^3 + E \left( \ell n F_v \right)^4 \right] \right\} \times 1.25 \]

Since \( V = \left( \frac{F}{F} \right) = 0.25 \times 1000 = 250 \text{ lbmol/h} \),
\[ W_v = V \left( \frac{M W_v}{V} \right) = 250 \left( \frac{26.15}{1} \right) = 6537.5 \text{ lb / h} \]
\[ L = F - V = 1000 - 250 = 750 \text{ lbmol/h}, \text{ and } W_L = (L) \left( \frac{M W_L}{L} \right) = (750)(20.82) = 15,615 \text{ lb/h}, \]
\[ F_v = W_L \sqrt{\frac{P_L}{P_V}} = \left( \frac{15615}{6537.5} \right) \frac{8.98 \times 10^{-4}}{.925} = 0.0744, \text{ and } \ell n (F_v) = -2.598 \]
\[ K_{\text{drum, vertical}} = 0.442, \text{ and } K_{\text{drum, horiz}} = 0.5525 \]
\[ u_{\text{perm}} = 0.5525 \sqrt{\frac{0.925 - 8.98 \times 10^{-4}}{8.98 \times 10^{-4}}} = 17.74 \text{ ft/s} \]
\[ A_{cs} = \frac{V \left( \frac{M W_v}{V} \right)}{u_{\text{perm}} 3600 \rho_v} = \frac{250 \left( 26.15 \right) \left( 454 \text{ g/lbm} \right)}{(17.74)(3600) \left( 8.98 \times 10^{-4} \text{ g/ml} \right) \left( 28316.85 \text{ ml/ft}^3 \right)} \]
\[ A_{cs} = 1.824 \text{ ft}^2, \quad A_T = A_{cs} / 0.2 = 9.12 \text{ ft}^2 \]

With \( L/D = 4, \quad D = \sqrt{4A_T/\pi} = 3.41 \text{ ft and } L = 13.6 \text{ ft} \)

2.D13. \textit{New Problem 4th ed.} \( x_{\text{butane}} = 1 - x_E = 0.912, y_{\text{butane}} = 1 - y_E = 0.454. K_E = y_E/x_E = 0.546/0.088 = 6.20, K_{\text{butane}} = y_B/x_B = 0.454/0.912 = 0.498. \)

Plot \( K_E \) and \( K_{\text{butane}} \) on DePriester chart. Draw straight line between them. Intersections with \( T \) and \( P \) axis give \( T_{\text{drum}} = 15 \text{ oC}, \text{ and } p_{\text{drum}} = 385 \text{ kPa from Figure 2-12.} \)

Use mass balances to find \( V/F: F = L + V \) and \( Fz_E = Lx_E + Vy_E. \) Substitute \( L = F - V \) into ethane balance and divide both sides by \( F. \) Obtain: \( z = (1 - V/F)x + y(V/F). \)

Solve for \( V/F = (z-x)/(y-x) = (0.36 - 0.088)/(0.546 - 0.088) = 0.594. \)

Spreadsheet used as a check (using \( T=15 \) and \( p = 385 \)) gave \( V/F = 0.593. \)

2.D14. \textit{New Problem 4th ed.} DePriester chart, Fig. 2-12: \( K_{C1} = 50, K_{C4} = 1.1, \text{ and } K_{C5} = 0.37; z_1 = 0.12, z_4 = 0.48, z_5 = 0.40 \)

Rachford-Rice equation:
\[ \frac{(K_{C2} - 1)z_{C1}}{1 + (K_{C1} - 1)\frac{V}{F}} + \frac{(K_{C4} - 1)z_{nC4}}{1 + (K_{nC4} - 1)\frac{V}{F}} + \frac{(K_{nC5} - 1)z_{nC5}}{1 + (K_{C5} - 1)\frac{V}{F}} = 0 \]

Equation becomes:
\[ 5.88 + 0.048 - 0.252 = 0 \]

Trials: \( V/F = 0.4, \text{ Eq. } = -0.005345; \text{ V/F = 0.39, Eq. } = 0.004506; \text{ V/F = 0.394, Eq. } = 0.000546, \) which is close enough with DePriester chart.

Liquid mole fractions:
\[ x_{C1} = \frac{z_{C1}}{1 - (K_{C1} - 1)(V/F)} = 0.12 \quad x_{C4} = 0.4618, x_{C5} = 0.5321, \text{ and } \sum x_i = 0.9998 \]

Vapor mole fractions: \( y_i = K_i x_i \): \( y_{C1} = 50(0.00591) = 0.2955, y_{C4} = 0.5080, y_{C5} = 0.1969, \sum y_i = 1.0004. \)
2.D15. This is an unusual way of stating problem. However, if we count specified variables we see that problem is not over or under specified. Usually V/F would be the variable, but here it isn’t. We can still write R-R eqn. Will have three variables: \( z_{C2} \), \( z_{iC4} \), \( z_{nC4} \). Need two other eqns: 
\[
\frac{z_{iC4}}{z_{nC4}} = \text{constant}, \quad \text{and} \quad z_{C2} + z_{iC4} + z_{nC4} = 1.0
\]
Thus, solve three equations and three unknowns simultaneously.

**Do It.** Rachford-Rice equation is,
\[
\frac{(K_{C2} - 1) z_{C2}}{1 + (K_{C2} - 1) V/F} + \frac{(K_{iC4} - 1) z_{iC4}}{1 + (K_{iC4} - 1) V/F} + \frac{(K_{nC4} - 1) z_{nC4}}{1 + (K_{nC4} - 1) V/F} = 0
\]
Can solve for \( z_{C2} = 1 - z_{iC4} \) and \( z_{iC4} = .8 z_{nC4} \). Thus \( z_{C2} = 1 - 1.8 z_{nC4} \)
Substitute for \( z_{iC4} \) and \( z_{C2} \) into R-R eqn.
\[
\frac{(K_{C2} - 1) (1-1.8 z_{nC4})}{1 + (K_{C2} - 1) V/F} + \frac{.8(K_{iC4} - 1) z_{nC4}}{1 + (K_{iC4} - 1) V/F} + \frac{(K_{nC4} - 1) z_{nC4}}{1 + (K_{nC4} - 1) V/F} = 0
\]

Thus,
\[
z_{nC4} = \frac{1.8 (K_{C2} - 1)}{1 + (K_{C2} - 1) V/F} = \frac{.8(K_{iC4} - 1)}{1 + (K_{iC4} - 1) V/F} = \frac{(K_{nC4} - 1)}{1 + (K_{nC4} - 1) V/F}
\]
Can now find K values and plug away. \( K_{C2} = 2.92, \ K_{iC4} = .375, \ K_{nC4} = .26 \)
Solution is \( z_{nC4} = 0.2957, \ z_{iC4} = .8 (.2957) = 0.2366, \) and \( z_{C2} = 0.4677 \)

2.D16. \( z_{c1} = 0.5, \ z_{c4} = 0.1, \ z_{c5} = 0.15, \ z_{c6} = 0.25, \ K_{c1} = 50, \ K_{c4} = .6, \ K_{c5} = .17, \ K_{c6} = 0.05 \)
1st guess. Can assume all \( C_1 \) in vapor, \( 1/3 C_4 \) in vapor, \( C_5 \) & \( C_6 \) in bottom
\[
\frac{V}{F}_1 = .5 + (.1)/3 = .53 \quad \text{This first guess is not critical.}
\]
R.R. eq. 
\[
\frac{V}{F} = \sum \frac{(K_i - 1) z_i}{1 + (K_i - 1) V/F} = 0
\]
\[
\frac{49(.5)}{1 + 49(.53)} + \frac{(-4)(.1)}{1-.4(.53)} + \frac{(-.83)(.15)}{1-.83(.53)} + \frac{(-.95)(.25)}{1-.95(.53)} = 0.157
\]

Eq. 3.33
\[
\frac{V}{F}_2 = \left( \frac{V}{F}_1 \right) + \frac{f \left( \frac{V}{F}_1 \right) \sum z_i (K_i - 1)^2}{\left[ 1 - (K_i - 1) \frac{V}{F} \right]^2}
\]
where \( (V/F)_1 = 0.53 \) and \( f (V/F)_1 = 0.157 \).

**calculate**
\[
(V/F)_2 = .53 + 0.157/2.92 = 0.584
\]
\[
V = .584 (150) = 87.6 \text{ kmol/h} \quad \text{and} \quad L = 150 - 87.6 = 62.4
\]
\[
x_{Cl} = \frac{z_{Cl}}{1 - (K_{Cl} - 1)(V/F)} = \frac{.5}{1 + 49(.584)} = 0.016883
\]
\[
y_{Cl} = K_{Cl} x_{Cl} = 50(0.016883) = 0.844
\]

Similar for other components.

2-D17.  

a.  \( V = 0.4F = 400, \quad L = 600 \)  
Slope = \(-L/F = -1.5\)  
Intercepts \( y = x = z = 0.70 \).  Plot line and find \( x_A = 0.65, \quad y_A = 0.77 \) (see graph)

b.  \( V = 2000, \quad L = 3000 \).  Rest identical to part a.

c.  Lowest \( x_A \) is horizontal op line (\( L = 0 \)).  \( x_A = 0.12 \)  
Highest \( y_A \) is vertical op line (\( V = 0 \)).  \( y_A = 0.52 \).  See graph

d.  \( V = 600, \quad L = 400, \quad -L/V = -0.667 \).
Find \( x_A = 0.40 \) on equilibrium curve.  Plot op line & find intersection point with \( y = x \) line.  \( z_A = 0.52 \)

2-D18.  

From \( x_i = \frac{z_i}{1 + \left( K_i - 1 \right) \frac{V}{F}} \), we obtain \( V = \frac{\frac{z_i}{x_i} - 1}{K_i - 1} \)

Guess \( T_{\text{drum}} \), calculate \( K_h, \quad K_b \) and \( K_p \), and then determine \( V/F \).

Check: \( \sum \frac{(K_i - 1)z_i}{1 + (K_i - 1)V/F} = 0 \) ?

Initial guess: \( T_{\text{drum}} \) must be less than temperature to boil pure hexane
\( K_h = 1.0, \ T = 94^\circ C \). Try 85°C as first guess (this is not very critical and the calculation will tell us if there is a mistake). \( K_h = 0.8, \ K_p = 4.8, \ K_p = 11.7 \).

\[
\frac{V}{F} = \frac{0.6}{0.85 - 1} = 1.471. \quad \text{Not possible. Must have} \quad K_h < \frac{0.6}{0.85} = 0.706.
\]

Try \( T = 73^\circ C \) where \( K_h = 0.6 \). Then \( K_h = 3.8, \ K_p = 9.9 \).

\[
\frac{V}{F} = \frac{0.6}{.6 - 1} = 0.735
\]

Check:

\[
\sum \frac{(K_i - 1)z_i}{1 + (K_i - 1)V/F} = \frac{(8.9)(1)}{1 + (8.9).735} + \frac{(2.8)(3)}{1 + (2.8).735} + \frac{(-4)(6)}{1 - (-4)(735)} = 0.05276
\]

Converge on \( T \sim 65.6^\circ C \) and \( V/F \sim 0.57 \).

2.D19. 90% recovery n-hexane means \((0.9)(Fz_{C6}) = L(x_{C6})\)

Substitute in \( L = F - V \) to obtain \( z_{C6}(.9) = (1 - V/F)x_{C6} \)

C8 balance: \( z_{C6}F = Lx_{C6} + Vy_{C6} = (F - V)x_{C6} + K_{C6}Vx_{C6} \)

or \( z_{C6} = (1 - V/F)x_{C6} + x_{C6}K_{C6}V/F \)

Two equations and two unknowns. Remove \( x_{C6} \) and solve

\[
z_{C6} = .93C_6 + \frac{(9)z_{C6}KV/F}{1 - V/F}
\]

Solve for \( V/F \):

\[
\frac{V}{F} = \frac{1}{(9K_{C6}) + .1}. \quad \text{Trial and error scheme.}
\]

Pick \( T \), Calc \( K_{C6} \), Calc \( V/F \), and Check \( f(V/F) = 0 \)?

If not \( K_{ref,new} = \frac{K_{ref}(T_{old})}{1 + df}(T) \)

Try \( T = 70^\circ C \). \( K_{C4} = 3.1, \ K_{C5} = .93, \ K_{C6} = .37 = K_{ref} \)

\[
\frac{V}{F} = \frac{.1}{(9)(.37) + .1} = 0.231.
\]

Rachford Rice equation

\[
f = \frac{(2.1).4}{1 + (2.1).231} + \frac{(-.08).25}{1 - (0.8).231} - \frac{(.63).35}{1 - (.63)(.231)} = .28719
\]

\[
K_{ref}(T_{new}) = \frac{.37}{1 + 0.28719} = 0.28745 \quad \text{(use .28)}
\]

Converge on \( T_{new} \sim 57^\circ C \). Then \( K_{C4} = 2.50, \ K_{C8} = .67, \) and \( V/F = 0.293 \).

2. D21. a.) \( K_{C2} = 4.8 \) \( K_{C5} = 0.153 \)

Soln to Binary R.R. eq. \[ \frac{V}{F} = \frac{-z_A}{(K_B - 1)} - \frac{z_B}{(K_A - 1)} \]
\[ \frac{V}{F} = \frac{-0.55}{(0.153 - 1)} - \frac{0.45}{(4.8 - 1)} = 0.5309 \]

\[ x_{c2} = \frac{z_{c2}}{1+(K_{c2}-1)\frac{V}{F}} = \frac{0.55}{1+(3.8)(0.5309)} = 0.1823, \quad y_{c2} = 0.8749, \quad x_{c5} = 0.8177, \quad y_{c5} = 0.1251 \]
Need to convert \( F \) to kmol. Avg MW = \( 0.55(30.07) + 0.45(72.15) = 49.17 \)

\[
F = 100,000 \frac{\text{kg}}{\text{hr}} \left( \frac{\text{kmol}}{49.17 \text{ kg}} \right) = 2033.7 \text{ kmol/h}, \quad V = \left( \frac{V}{F} \right) F = 1079.7, \quad L = F - V = 954.0 \text{ kmol/h}
\]

b.) \( u_{\text{perm}} = K_{\text{drum}} \sqrt{\frac{\rho_L}{\rho_v}} \)

To find

\[
MW_L = (0.1823)(30.07) + (0.8177)(72.15) = 64.48
\]

\[
MW_v = (0.8749)(30.07) + (0.1251)(72.15) = 35.33
\]

For liquid assume ideal mixture:

\[
\bar{V}_L = x_{C2} \bar{V}_{C2,\text{liq}} + x_{C5} \bar{V}_{C5,\text{liq}} = x_{C2} \frac{MW_{C2}}{\rho_{C2,\text{liq}}} + x_{C5} \frac{MW_{C5}}{\rho_{C5,\text{liq}}}
\]

\[
\bar{V}_L = (0.1823)\left( \frac{30.07}{0.54} \right) + (0.8177)\left( \frac{72.15}{0.63} \right) = 103.797 \text{ ml/mol}
\]

\[
\rho_L = \frac{MW_L}{\bar{V}_L} = \frac{64.48}{103.797} = 0.621 \text{ g/ml}
\]

For vapor: ideal gas:

\[
\rho_v = \frac{MW_v}{RT} = \frac{700 \text{ kPa}}{101.3 \text{ kPa}} \left( \frac{35.33 \text{ g}}{\text{mol}} \right) \frac{82.0575 \text{ ml atm}}{\text{mol K}} (303.16 \text{K}) = 0.009814 \text{ g/ml}
\]

\[K_{\text{drum}} : \text{Use Eq. (2-60) with } F_{lv} = \frac{W_l}{W_v} \sqrt{\frac{\rho_v}{\rho_L}} \]

\[
W_L = 997.7 \frac{\text{kmol}}{\text{h}} \left| \begin{array}{c} 64.48 \text{ kg} \\ \text{kmol} \end{array} \right| = 6,4331.7 \text{ kg/h}, \quad W_v = 881.5[35.33] = 31,143.4 \text{ kg/h}
\]

\[
F_{lv} = \frac{64331.7}{31,143.3} \sqrt{0.009814} = 0.2597
\]

\[K_{\text{drum}} = \exp\left[ -1.877478 + (-0.81458)(\ln 0.2597) + (-0.18707)(\ln 0.2597)^2 \right.\]

\[\left. + (-0.0145229)(\ln 0.2597)^3 + (-0.0010149)(\ln 0.2597)^4 \right] = 0.3372
\]

\[
u_{\text{perm}} = (0.3372) \sqrt{\frac{0.621 - 0.009814}{0.009814}} \left( \frac{1.0 \text{ m}}{3.2808 \text{ ft}} \right) = 0.8111 \text{ m/s}
\]

\[
A_c = \frac{V MW_v}{3600 \rho_v} = \frac{1079.7 \text{ kmol/h}}{3600 \text{ s}} \left( \frac{10^6 \text{ cm}^3}{\text{m}^3} \right) = 1.392 \text{ m}^2
\]

\[
D = \sqrt{4A_c / \pi} = 1.33 \text{ m}. \quad \text{Arbitrarily} \quad L/D = 4, \quad L = 5.32 \text{ m}
\]


a. \( V = F - L = 50 - 20 = 30 \text{ kmol/h} \). \( V/F = 3/5 \), Slope operating line = \(-L/V = -20/30 = -2/3 \), \( z_M = 0.7 \)
From graph, \( y = 0.8, \ x = 0.54 \).

b. From graph of \( T \) vs. \( x_M \), \( T_{	ext{drum}} = 72.3^\circ \text{C} \). (see graph).


Part a. \( F_{\text{new}} = (1500 \ \text{kmol/h})(1.0 \ \text{lbmol}/(0.45359 \ \text{kmol})) = 3307 \ \text{lb mol/h} \).

\( V, \ W_V, \ L, \) and \( W_L \) are the values in Example 2-4 divided by 0.45359. The conversion factor divides out in \( F_{lv} \) term. Thus, \( F_{lv}, \ K_{\text{drum}}, \) and \( u_{\text{perm}} \) are the same as in Example 2-4. The Area increases because \( V \) increases: \[ \text{Area} = \frac{\text{Area}_{	ext{Example 2-4}}}{0.45359} = \frac{16.047}{0.45359} = 35.38 \ \text{ft}^2. \]

\[ \text{Diameter} = \sqrt{4 \cdot \text{Area} / \pi} = 6.71 \ \text{feet} \]

Probably round this off to 7.0 feet and use a drum height of 28 feet.

b. \( F_{\text{parallel}} = 3307 - 1500 = 1807 \ \text{lbmol/h}. \)

\( F_{lv}, \ K_{\text{drum}}, \) and \( u_{\text{perm}} \) are the same as in Example 2-4. \[ V_{\text{parallel}} = \left(\frac{V}{F}\right) F_{\text{parallel}} = 0.51 \times 1807 = 921.6 \ \text{kmol/h}. \]
\[ A = 16.047 \times \frac{V_{\text{parallel}}}{V} = 16.047 \times (921.6 / 765) = 19.33 \text{ ft}^2 \]

Then, \( \text{Diameter} = \sqrt{4 \times A / \pi} = 4.96 \text{ feet} \), Use a 5.0 feet diameter and a length of 20 feet.

2.D24. \( p = 300 \text{ kPa} \) At any \( T \). \( K_{C3} = y_{C3}/x_{C3} \), \( K \)'s are known. \( K_{C6} = y_{C6}/x_{C6} = (1-y_{C3})/(1-x_{C3}) \)

Substitute 1\(^{st}\) equation into 2\(^{nd}\) \( K_{C6} = (1-K_{C3}x_{C3})/(1-x_{C3}) \)

Solve for \( x_{C3}, \) \[ x_{C3} = \frac{1-K_{C6}}{K_{C3}-K_{C6}} \] & \[ y_{C3} = \frac{K_{C3}(1-K_{C6})}{K_{C3}-K_{C6}} \]

At 300 kPa pure propane \( (K_{C3} = 1.0) \) boils at -14°C (Fig. 2-10)

At 300 kPa pure n-hexane \( (K_{C6} = 1.0) \) boils at 110°C

Check: at -14°C \( x_{C3} = \frac{1-K_{C6}}{1-K_{C6}} = 1, \) \( y_{C3} = \frac{1(1-K_{C6})}{1-K_{C6}} = 1.0 \)

at 110°C \( x_{C3} = 0, \) \( y_{C3} = \frac{K_{C3}(0)}{K_{C3}} = 0 \)

Pick intermediate temperatures, find \( K_{C3} \) & \( K_{C6} \), calculate \( x_{C3} \) & \( y_{C3} \).

<table>
<thead>
<tr>
<th>( T )</th>
<th>( K_{C3} )</th>
<th>( K_{C6} )</th>
<th>( x_{C3} )</th>
<th>( y_{C3} = K_{C3}x_{C3} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>0°C</td>
<td>1.45</td>
<td>0.027</td>
<td>( \frac{1-0.027}{1.45-0.027} = 0.684 )</td>
<td>0.9915</td>
</tr>
<tr>
<td>10°C</td>
<td>2.1</td>
<td>0.044</td>
<td>0.465</td>
<td>0.976</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>See</td>
</tr>
<tr>
<td>20°C</td>
<td>2.6</td>
<td>0.069</td>
<td>0.368</td>
<td>0.956</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Graph</td>
</tr>
<tr>
<td>30°C</td>
<td>3.3</td>
<td>0.105</td>
<td>0.280</td>
<td>0.924</td>
</tr>
<tr>
<td>40°C</td>
<td>3.9</td>
<td>0.15</td>
<td>0.227</td>
<td>0.884</td>
</tr>
<tr>
<td>50°C</td>
<td>4.7</td>
<td>0.21</td>
<td>0.176</td>
<td>0.827</td>
</tr>
<tr>
<td>60°C</td>
<td>5.5</td>
<td>0.29</td>
<td>0.136</td>
<td>0.75</td>
</tr>
<tr>
<td>70°C</td>
<td>6.4</td>
<td>0.38</td>
<td>0.103</td>
<td>0.659</td>
</tr>
</tbody>
</table>

b. \( x_{C3} = 0.3 \), \( V/F = 0.4 \), \( L/V = 0.6/0.4 = 1.5 \)

Operating line intersects \( y = x = 0.3 \), Slope -1.5

\[ y = \frac{-L}{V}x + \frac{F}{V}z \] at \( x = 0, \) \( y = \frac{F}{V}z = \frac{0.3}{0.4} = 0.75 \)

Find \( y_{C3} = 0.63 \) and \( x_{C3} = 0.062 \)

Check with operating line: \( 0.63 = -1.5(0.062) + 0.75 = 0.657 \) OK within accuracy of the graph.

c. Drum \( T \): \( K_{C3} = y_{C3}/x_{C3} = 0.63/0.062 = 10.2 \), DePriester Chart \( T = 109^\circ C \)

d. \( y = .8, \) \( x \sim .16 \) Slope = \( \frac{\Delta y}{\Delta x} = \frac{.8-.6}{.16-.6} = -0.45 = \frac{-1-f}{f} = -.45 \)

\( V/F = f = 1/1.45 = 0.69 \)
2.D25.  20% Methane and 80% n-butane. \( T_{\text{drum}} = .50 \, ^\circ\text{C} \), \( \frac{V}{F} = 0.40 \), Find \( p_{\text{drum}} \)

\[
0 = f\left( \frac{V}{F} \right) = \frac{(K_A-1)z_A}{1+(K_A-1)\frac{V}{F}} + \frac{(K_B-1)z_B}{1+(K_B-1)\frac{V}{F}}
\]

---

Pick \( p_{\text{drum}} = 1500 \) kPa: \( K_{C4} = 13 \), \( K_{nc4} = 0.4 \)

(Any pressure with \( K_{C1} > 1 \) and \( K_{C4} < 1.0 \) is OK)

Trial 1

\[
f_i = \frac{12.2}{1+12.4} + \frac{(-.6)(.8)}{1-.6(.4)} = -0.2178 \quad \text{Need lower} \ p_{\text{drum}}
\]

\[
V_0f = \frac{0.4}{1+(-.2178)} = 0.511 \quad \text{with} \ d = 1.0
\]

\[
P_{\text{new}} = 1160 \quad K_{C1} = 16.5, f_2 = \frac{(15.5)(.2)}{1+15.5(.4)} + \frac{-489(.8)}{1-(.489)(.4)} = 0.4305 + -.4863 = -0.05769
\]

\[
K_{C4}(P_{\text{new}}) = \frac{K_{C4}(P_{\text{old}})}{1+(d)f(P_{\text{old}})} = \frac{(0.4)}{1+(-.2178)} = 0.511 \quad \text{with} \ d = 1.0
\]

\[
P_{\text{new}} = 1160 \quad K_{C1} = 16.5, f_2 = \frac{(15.5)(.2)}{1+15.5(.4)} + \frac{-489(.8)}{1-(.489)(.4)} = 0.4305 + -.4863 = -0.05769
\]

\[
K_{C4}(P_{\text{new}}) = \frac{0.511}{1+0.05769} = 0.541, \quad p_{\text{new}} = 1100, \quad K_{C1} = 17.4
\]

\[
f_3 = \frac{16.4(.2)}{1+(16.4)(.4)} + \frac{(-.459)(.8)}{1-(.459)(.4)} = -0.0159, \quad \text{OK.} \quad \text{Drum pressure} = 1100 \, \text{kPa}
\]

\[
x_i = \frac{z_i}{1+(K_i-1)\frac{V}{F}}, \quad x_{C1} = \frac{0.2}{1+(16.4)(.4)} = 0.02645
\]

\[
y_{C1} = K_{C1}x_{C1} = (17.4)(0.02645) = 0.4603
\]

2.D26. a) Can solve for \( L \) and \( V \) from M.B. \( 100 = F = V + L \)

\[45 = Fz = 0.8V + 0.2162L\]

Find:

\[L = 59.95 \quad \text{and} \quad V = 40.05\]

b) Stage is equil. \( K_{C3} = \frac{y_3}{x_3} = \frac{0.8}{0.2162} = 3.700 \), \( K_{C5} = \frac{0.2}{0.7838} = 0.2552 \)

These \( K \) values are at same \( T, P \). Find these 2 \( K \) values on DePriester chart. Draw straight line between them. Extend to \( T_{\text{drum}}, \ p_{\text{drum}} \). Find \( 10^\circ\text{C}, \ 160 \) kPa.

2.D27. a) \( VP_{C5} : \log_{10} VP = 6.853 - \frac{1064.8}{0+233.01} = 2.2832, \quad VP = 191.97 \, \text{mmHg}\)

b) \( VP = 3 \times 760 = 2280 \, \text{mmHg}, \quad \log_{10} VP = (6.853) - (1064.8)/(T + 233.01)\)

Solve for \( T = 71.65^\circ\text{C}\)

c) \( P_{\text{tot}} = 191.97 \, \text{mm Hg} \) [at boiling for pure component \( P_{\text{tot}} = VP \)]

d) \( C5: \log_{10} VP = 6.853 - \frac{1064.8}{30+233.01} = 2.8045, \quad VP = 637.51 \, \text{mm Hg} \)

\[K_{C5} = \frac{VP_{C5}}{P_{\text{tot}}} = \frac{637.51}{500} = 1.2750\]
C6: \( \log_{10} VP_{C6} = 6.876 - \frac{1171.17}{30 + 224.41} = 2.2725 \), \( VP_{C6} = 187.29 \text{ mm Hg} \)

\( K_{C6} = \frac{187.29}{500} = 0.3746 \)

e.) \( K_A = \frac{y_A}{x_A} \quad K_B = \frac{y_B}{x_B} = \frac{1-y_A}{1-x_A} \)

If \( K_A \) & \( K_B \) are known, two eqns. with 2 unknowns \( (K_A \) & \( y_A \) \) Solve.

\[
\begin{align*}
C_5 &: \quad \frac{1-K_{C6}}{K_{C5}-K_{C6}} = \frac{1-0.3746}{1.2750-0.3746} = 0.6946 \\
C_5 &= (1.2750)(0.6946) = 0.8856
\end{align*}
\]

f.) Overall, \( M.B., F = L + V \) or \( 1 = L + V \)

\( C_5 : \quad Fx_L + Vy_5 = \frac{.75}{0.6946} L + \frac{0.8856}{0.6946} V \)

Solve for \( L \) & \( V \): \( L = 0.7099 \) & \( V = 0.2901 \text{ mol} \)

\( g.) \) Same as part \( f \), except units are \( \text{mol/min} \).


From example 2-4, \( x_H = 0.19, T_{drum} = 378K, V/F = 0.51, y_H = 0.6, z_H = 0.40 \)

\( \text{MW}_v = 97.39 \text{ lbm/lbmole (Example 2-4)} \)

\[
\rho_v = 3.14 \times 10^{-3} \text{ g/mol} \left( \frac{1}{454\text{g/lbm}} \right) \left( \frac{28316.85\text{cm}^3}{\text{ft}^3} \right) = 0.198 \text{ lbm/ft}^3
\]

Example 2.4

\( u_{perm} = K_{drum} \sqrt{\frac{\rho_L - \rho_v}{\rho_v}}, \quad K_{horiz} = 1.25 K_{vertical} \)

From Example 2-4, \( K_{vertical} = 0.4433, K_{horiz} = 1.25(0.4433) = 0.5541 \)

\[ u_{perm} = 0.5541 \left( \frac{0.6960 - 0.00314}{0.00314} \right)^{1/2} = 8.231 \text{ ft/s} \] [densities from Example 2-4]
42

\[ V = \left( \frac{V}{F} \right) F = (0.51) \left( 3000 \frac{\text{lbmol}}{\text{h}} \right) = 1530 \ \text{lbmol/h} \]

\[ A_{\text{vap}} = \frac{1530 \frac{\text{lbmol}}{\text{h}} \left( 97.39 \frac{\text{lb}}{\text{lbmol}} \right)}{8.231 \frac{\text{ft}}{\text{s}}} \left( 3600 \frac{\text{s}}{\text{h}} \right) \left( 0.1958 \frac{\text{lbm}}{\text{ft}^3} \right) = 25.68 \ \text{ft}^2 \]

\[ A_{\text{total}} = A_{\text{vap}} / 0.2 = 128.4 \text{ft}^3, D_{\min} = \sqrt{4 A_{\text{total}} / \pi} = 12.8 \text{ft} \]

\[ V_{\text{liq}} = \frac{160.068}{43.41} \frac{55+85}{60 \text{ min/ h}} = 8603.8 \text{ft}^3, h = \frac{5 V_{\text{liq}}}{\pi D^2} = 83.51 \text{ft} \text{ and } h/D = 6.5. \]

2.D29. The stream tables in Aspen Plus include a line stating the fraction vapor in a given stream. Change the feed pressure until the feed stream is all liquid (fraction vapor = 0). For the Peng-Robinson correlation the appropriate pressure is 74 atm.

The feed mole fractions are: methane = 0.4569, propane = 0.3087, n-butane = 0.1441, i-butane = 0.0661, and n-pentane = 0.0242.

b. At 74 atm, the Aspen Plus results are: L = 10169.84 kg/h = 201.636 kmol/h, V = 4830.16 kg/h = 228.098 kmol/h, and T_{drum} = -40.22 °C.

The vapor mole fractions are: methane = 0.8296, propane = 0.1458, n-butane = 0.0143, i-butane = 0.0097, and n-pentane = 0.0006.

The liquid mole fractions are: methane = 0.0353, propane = 0.4930, n-butane = 0.2910, i-butane = 0.1298, and n-pentane = 0.0509.

c. Aspen Plus gives the liquid density = 0.60786 g/cc, liquid avg MW = 50.4367, vapor density = 0.004578 g/cc = 4.578 kg/m³, and vapor avg MW = 21.17579 g/mol = kg/kmol.

The value of uperm (in ft/s) can be determined by combining Eqs. (2-64), (2-65) and (2-69)

\[ F_{lv} = (W_l/W_v)\left[\frac{\rho_v}{\rho_l}\right]^{0.5} = (10169.84/4830.16)[0.004578/0.60786]^{0.5} = 0.18272 \]

Resulting K_{vertical} = 0.378887, K_{horizontal} = 0.473608, and u_{perm} = 5.436779 ft/s = 1.657 m/s

\[ A_{\text{vap}} = \frac{4830.16 \frac{\text{kg}}{\text{h}}}{\left( 1.657 \frac{\text{m}}{\text{s}} \right) \left( 3600 \frac{\text{s}}{\text{h}} \right) \left( 4.578 \frac{\text{kg}}{\text{m}^3} \right)} = 0.177 \text{ m}^2 \]

\[ A_{\text{total}} = A_{\text{vap}} / 0.2 = 0.884 \text{ m}^3, D_{\min} = \sqrt{4 A_{\text{total}} / \pi} = 1.06 \text{ m} \]

\[ h / D = 6 = \frac{5 V_{\text{liq}}}{\pi D^2}, \text{ thus } V_{\text{liq}} = 6 \pi D^2 / 5 = 4.23 \text{ m}^3 \]

\[ V_{\text{liq}} = (\text{Vol rate})(\text{hold time + surge time}) = \left( \frac{10169.84 \text{ kg}}{\text{h}} \right) \left( \frac{10169.84 - 9}{60} + \text{st} \right) \]

\[ \text{st} = 607.86 V_{\text{liq}} / 10169.84 - 9 / 60 = 0.103 \text{ hours} = 6.18 \text{ min} \]

2.D30. a. From the equilibrium data if y_A = .40 mole fraction water, then x_A = 0.09 mole fraction water. Can find L_A and V_A by solving the two mass balances for stage A simultaneously.

L_A + V_A = F_A = 100 and L_A (.09) + V_A (.40) = (100) (.20). The results are V_A = 35.48 and L_A = 64.52.
b. In chamber B, since 40 % of the vapor is condensed, \((V/F)_B = 0.6\). The operating line for this flash chamber is,
\[
y = -(L/V)x + F_B/V \left( z_B \right)
\]
where \(z_B = y_A = 0.4\) and \(L/V + .4F_B/.6F_B = 2/3\). This operating line goes through the point \(y = x = z_B = 0.4\) with a slope of \(-2/3\). This is shown on the graph. Obtain \(x_B = 0.18\) & \(y_B = 0.54\).

\[L_B = \text{(fraction condensed)(feed to B)} = 0.4(35.48) = 14.19 \text{ kmol/h} \text{ and } V_B = F_B - L_B = 21.29.\]

c. From the equilibrium if \(x_B = 0.20, y_B = 0.57\). Then solving the mass balances in the same way as for part a with \(F_B = 35.48\) and \(z_B = 0.4\), \(L_B = 16.30\) and \(V_B = 19.18\). Because \(x_B = z_A\), recycling \(L_B\) does not change \(y_B = 0.57\) or \(x_A = 0.09\), but it changes the flow rates \(V_{B,new}\) and \(L_{A,new}\). With recycle these can be found from the overall mass balances: \(F = V_{B,new} + L_{A,new}\) and \(Fz_A = V_{B,new}y_B + L_{A,new}x_A\). Then \(V_{B,new} = 22.92\) and \(L_{A,new} = 77.08\).

Graph for problem 2.D30.


a) Since K’s are for mole fractions, need to convert feed to mole fractions.

<table>
<thead>
<tr>
<th>Basis: 100 kg feed</th>
<th>50 kg n C_4 1 kmol 58.12 kg = 0.8603 kmol</th>
<th>(z_4 = 0.555)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>50 kg n C_5 1 kmol 72.5 kg = 0.6897 kmol n C_5</td>
<td>(z_5 = 0.445)</td>
</tr>
<tr>
<td></td>
<td>Total 1.5499 kmol</td>
<td></td>
</tr>
</tbody>
</table>
DePriester Chart \( K_{C4} = 2.05, \) \( K_{C5} = 0.58, \) (Result similar if use Raoult's law).

\[
\frac{V}{F} = \frac{-0.555 - 0.445}{0.58 - 1} = 1.3214 - 0.424 = 0.8976
\]

Check \( f \left( \frac{V}{F} \right) = \frac{\left(1.05\right)\left(5.55\right)}{1+\left(5.55\right)} + \frac{\left(-0.42\right)\left(4.45\right)}{1-\left(0.42\right)} = 0.3000 - 0.29999 = 0 \) OK

Eq. 3.23 \( x_{c4} = \frac{z_{c4}}{1+\left(K_{c4} - 1\right)V/F} = \frac{0.555}{1+1.056(0.8976)} = 0.2857 \),

\( x_{c5} = \frac{7.143}{y_{c4} = \frac{K_{c4}}{x_{c4}} = 0.5857, \ y_{c5} = 0.4143} \)

b) From problem 2.D.g., \( K_{C4} = 1.019 \) and \( K_{C5} = 0.253 \).

Solving RR equation,

\[
\frac{V}{F} = \left[ -\frac{z_A}{\left(K_B - 1\right)} - \frac{z_B}{\left(K_A - 1\right)} \right] = -\frac{0.555}{(0.253 - 1)} - \frac{0.445}{0.019} = -23.28
\]

NOT possible. Won't flash at 0ºC.


\( (V/F)_A = 2/3, \ L/V = \frac{1}{3} = 1/2 \) Slope = \( -1/2 \) Through \( y = x = z_A = 0.6 \) See figure

a. \( L_A = \frac{1}{3}F = 33.33, \ x_{MA} = 0.375 \) (from Figure) \( V_A = \frac{2}{3}F = 66.67, \ y_{MA} = 0.72 \) (from Figure)

\[
\frac{V}{F_B} = 0.4 - \left( \frac{L}{V_B} \right) = -1.5
\]

Through \( y = x = z_B = y_A = 0.72 \)

\( V_B = 0.4F_B = 0.4V_A = 0.4(66.67) = 26.67 \), \( L_B = 0.6F_B = 0.6(66.67) = 40.00 \)

b. \( z_C = x_A = 0.375, \ x_C = 0.15, \ F_C = L_A = 33.33 \), From equilibrium \( y_C = 0.51 \)

At \( x = 0, \ y_C = 0.60 = \left( \frac{F}{V} \right) z_C \Rightarrow \left( \frac{V}{F} \right) = \frac{z_C}{y_C} = \frac{0.375}{0.6} = 0.625 \)

\( V_C = \left( \frac{V}{F} \right) F_C = 0.625(33.33) = 20.83, \ L_C = F_C - V_C = 33.33 - 20.83 = 12.5 \)
2. E1. From Aspen Plus run with 1000 kmol/h at 1 bar, \( L = V = 500 \) kmol/h, \( W_L = 9212.78 \) kg/h, \( W_V = 13010.57 \) kg/h, liquid density = \( 916.14 \) kg/m\(^3\), liquid avg MW = 18.43, vapor density = 0.85 kg/m\(^3\), and vapor avg MW = \( 26.02 \), \( T_{drum} = 94.1 \) °C, and \( Q = 6240.85 \) kW.

The diameter of the vertical drum in meters (with \( u_{perm} \) in ft/s) is

\[
D = \left\{\frac{4(MW_V) V}{[3600 \pi \rho_v u_{perm} (1 \text{ m/3.281 ft})]}\right\}^{0.5} = \left\{\frac{4(26.02)(500)}{[3600(3.14159)(0.85)(1/3.281)u_{perm}]}\right\}^{0.5}
\]

\[
F_{lv} = \left(\frac{W_L}{W_V}\right)\left[\frac{\rho_V}{\rho_L}\right] = \left(\frac{9212.78}{13010.57}\right)\left[\frac{0.85}{916.14}\right]^{0.5} = 0.02157
\]

Resulting \( K_{vertical} = 0.404299 \), and \( u_{perm} = 13.2699 \) ft/s, and \( D = 1.16 \) m. Appropriate standard size would be used. Mole fractions isopropanol: liquid = 0.00975, vapor = 0.1903

b. Ran with feed at 9 bar and \( p_{drum} = 8.9 \) bar with \( V/F = 0.5 \). Obtain \( W_L = 9155.07 \) kg/h, \( W_V = 13068.27 \), density liquid = \( 836.89 \), density vapor = \( 6.37 \) kg/m\(^3\)

\[
D = \left\{\frac{4(MW_V) V}{[3600 \pi \rho_v u_{perm} (1 \text{ m/3.281 ft})]}\right\}^{0.5} = \left\{\frac{4(26.14)(500)}{[3600(3.14159)(6.37)(1/3.281)u_{perm}]}\right\}^{0.5}
\]

\[
F_{lv} = \left(\frac{W_L}{W_V}\right)\left[\frac{\rho_V}{\rho_L}\right]^{0.5} = \left(\frac{9155.07}{13068.27}\right)\left[\frac{6.37}{836.89}\right]^{0.5} = 0.06112
\]

Resulting \( K_{vertical} = .446199 \), \( u_{perm} = 5.094885 \) ft/s, and \( D = 0.684 \) m. Thus, the method is feasible.

c. Finding a pressure to match the diameter of the existing drum is trial and error. If we do a linear interpolation between the two simulations to find a pressure that will give us \( D = 1.0 \) m (if linear), we find \( p = 3.66 \). Running this simulation we obtain, \( W_L = 9173.91 \) kg/h, \( W_V = 13049.43 \), density liquid = \( 874.58 \), density vapor = \( 2.83 \) kg/m\(^3\), \( MW_v = 26.10 \)

\[
D = \left\{\frac{4(MW_V) V}{[3600 \pi \rho_v u_{perm} (1 \text{ m/3.281 ft})]}\right\}^{0.5} = \left\{\frac{4(26.10)(500)}{[3600(3.14159)(2.83)(1/3.281)u_{perm}]}\right\}^{0.5}
\]

\[
F_{lv} = \left(\frac{W_L}{W_V}\right)\left[\frac{\rho_V}{\rho_L}\right]^{0.5} = \left(\frac{9173.91}{13049.43}\right)\left[\frac{2.83}{874.58}\right]^{0.5} = 0.0400
\]

Resulting \( K_{vertical} = .441162 \), \( u_{perm} = 7.742851 \) ft/s, and \( D = 0.831 \) m.

Plotting the curve of \( D \) versus \( p_{drum} \) and setting \( D = 1.0 \), we interpolate \( p_{drum} = 2.1 \) bar At \( p_{drum} = 2.1 \) bar simulation gives, \( W_L = 9188.82 \) kg/h, \( W_V = 13034.53 \), density liquid = \( 893.99 \), density vapor = \( 1.69 \) kg/m\(^3\), \( MW_v = 26.07 \)

\[
D = \left\{\frac{4(MW_V) V}{[3600 \pi \rho_v u_{perm} (1 \text{ m/3.281 ft})]}\right\}^{0.5} = \left\{\frac{4(26.07)(500)}{[3600(3.14159)(1.69)(1/3.281)u_{perm}]}\right\}^{0.5}
\]

\[
F_{lv} = \left(\frac{W_L}{W_V}\right)\left[\frac{\rho_V}{\rho_L}\right]^{0.5} = \left(\frac{9188.82}{13034.53}\right)\left[\frac{1.69}{893.99}\right]^{0.5} = 0.0307
\]

Resulting \( K_{vertical} = .42933 \), \( u_{perm} = 9.865175 \) ft/s, and \( D = 0.953 \) m.

This is reasonably close and will work OK. \( T_{drum} = 115.42 \) °C, \( Q = 6630.39 \) kW, Mole fractions isopropanol: liquid = 0.00861, vapor = 0.1914

In this case there is an advantage operating at a somewhat elevated pressure.
2.E2. This problem was 2.D13 in the 2nd edition of SPE.

a. Will show graphical solution as a binary flash distillation. Can also use R-R equation. To generate equil. data can use

\[ x_{C6} + x_{C8} = 1.0, \text{ and } y_{C6} + y_{C8} = K_{C6} x_{C6} + K_{C8} x_{C8} = 1.0 \]

Substitute for \( x_{C6} \)

\[ x_{C6} = \frac{1 - K_{C8}}{K_{C6} - K_{C8}} \]

Pick \( T \), find \( K_{C6} \) and \( K_{C8} \) (e.g. from DePriester charts), solve for \( x_{C6} \). Then \( y_{C6} = K_{C6} x_{C6} \)

<table>
<thead>
<tr>
<th>( T^\circ C )</th>
<th>( K_{C6} )</th>
<th>( K_{C8} )</th>
<th>( x_{C6} )</th>
<th>( y_{C6} = K_{C6} x_{C6} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>125</td>
<td>4</td>
<td>1.0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>120</td>
<td>3.7</td>
<td>.90</td>
<td>.0357</td>
<td>.321</td>
</tr>
<tr>
<td>110</td>
<td>3.0</td>
<td>.68</td>
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<td>.141</td>
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<tr>
<td>100</td>
<td>2.37</td>
<td>.52</td>
<td>.2595</td>
<td>.615</td>
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<tr>
<td>90</td>
<td>1.8</td>
<td>.37</td>
<td>.4406</td>
<td>.793</td>
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<tr>
<td>80</td>
<td>1.4</td>
<td>.26</td>
<td>.650</td>
<td>.909</td>
</tr>
<tr>
<td>66.5</td>
<td>1.0</td>
<td>.17</td>
<td>1.0</td>
<td>1.0</td>
</tr>
</tbody>
</table>

Op Line Slope = \( \frac{L}{V} = \frac{1 - V/F}{V/F} = \frac{.6}{.4} = -1.5 \), Intersection \( y = x = z = 0.65 \).

See Figure. \( y_{C6} = 0.85 \) and \( x_{C6} = 0.52 \). Thus \( K_{C6} = .85/.52 = 1.63 \).
This corresponds to \( T = 86^\circ C = 359K \)

b. Follows Example 2-4.

\[ MW_L = x_{C6} (MW)_{C6} + x_{C8} (MW)_{C8} = (.52)(86.17) + (.48)(114.22) = 99.63 \]

\[ \bar{V}_L = x_{C6} \frac{(MW)_{C6}}{\rho_{C6}} + x_{C8} \frac{(MW)_{C8}}{\rho_{C8}} = (.52) \frac{86.17}{.659} + (.48) \frac{114.22}{.703} = 145.98 \text{ ml/mol} \]

\[ \rho_L = \frac{MW_L}{\bar{V}_L} = \frac{99.63}{145.98} = 0.682 \text{ g/ml} \left[ \frac{28316 \text{ ml/ft}^3}{454 \text{ g/lbm}} \right] = 42.57 \frac{\text{lbm}}{\text{ft}^3} \]
\[
\overline{\text{MW}_v} = y_{C_6} (\text{MW}_{C_6}) + y_{C_8} (\text{MW}_{C_8}) = 0.85(86.17) + 0.15(114.22) = 90.38
\]
\[
\rho_v = \frac{p\overline{\text{MW}_v}}{RT} = \frac{(1.0) 90.38 \text{ g/mol}}{82.0575 \text{ ml atm mol}^{-1} \text{K}^{-1}} = 0.00307 \text{ g/ml} = 0.19135 \text{ lbm/ft}^{-3}
\]

Now we can determine flow rates
\[
V = \left(\frac{V}{F}\right) F = (0.4)(10,000) = 4000 \text{ lbmol/h}
\]
\[
W_v = V \left(\overline{\text{MW}_v}\right) = 4000(90.38) = 361,520 \text{ lb/h}
\]

\[
L = F - V = 6000 \text{ lbmol/h}, \quad W_L = L \left(\overline{\text{MW}_L}\right) = (6000)(99.63) = 597,780 \text{ lb/h}
\]
\[
F_v = \frac{W_L}{W_v \sqrt{\rho_L}} = \frac{597,780}{361,520 \sqrt{42.57}} = 0.111, \quad \ell n F_v = -2.1995
\]
\[
K_{\text{drum}} = \exp \left[(-1.87748) + (-0.81458)(-2.1995) + (-0.18707)(-2.1995)^2
\right.
\]
\[
\left. + (-0.01452)(-2.1995)^3 + (-0.00101)(-2.1995)^4 \right] = 0.423
\]
\[
u_{\text{Perm}} = K_{\text{drum}} \sqrt{\rho_L - \rho_v / \rho_v} = (0.423) \sqrt{(42.57 - 19135)}/19135 = 6.30 \text{ ft/s}
\]
\[
A_{C_5} = \frac{V \left(\overline{\text{MW}_v}\right)}{u_{\text{Perm}} (3600) \rho_v} = \frac{(4000)(90.38)}{u_{\text{Perm}} (3600)(0.19135)} = 83.33 \text{ ft}^2
\]
\[
D = \sqrt{4A_{C_5}/\pi} = \sqrt{4(83.33)/\pi} = 10.3 \text{ ft}. \quad \text{Use 10.5 ft.}
\]

\[
L \text{ ranges from } 3 \times 10.5 = 31.5 \text{ ft to } 5 \times 10.5 = 52.5 \text{ ft.}
\]

Note: This \( u_{\text{Perm}} \) is at 85% of flood. If we want to operate at lower % flood (say 75%)
\[
u_{\text{Perm},75} = (0.75/0.85) u_{\text{Perm},85} = (0.75/0.85)(6.3) = 5.56
\]
Then at 75% of flood, \( A_{C_5} = 94.44 \) which is \( D = 10.96 \) or 11.0 ft.

2.E3. New problem 4th edition. The difficulty of this problem is it is stated in weight units, but the VLE data is in molar units. The easiest solution path is to work in weight units, which requires converting some of the equilibrium data to weight units and replotting – good practice. The difficulty with trying to work in molar units is the ratio \( L/V = 0.35/0.65 = 0.5385 \) in weight units becomes in molar units, \[
\frac{L_{\text{molar}}}{V_{\text{molar}}} = \frac{L_{\text{wt}} (MW)_{\text{vapor}}}{V_{\text{wt}} (MW)_{\text{liquid}}}, \quad \text{but } x \text{ and } y \text{ are not known the molecular weights are unknown.}
\]

In weight units, \( V = F(V/F) = 2000 \text{ kg/h} \times (0.35) = 700 \text{ kg/h}. \quad L = F - V = 1300 \text{ kg/h.}
\]

In weight units the equilibrium data (Table 2-7) can be converted as follows:

Basis: 1 mol, \( x = 0.4 \) and \( y = 0.729, T = 75.3 \text{ C}

Liquid: 0.4 mol methanol \( \times 32.04 \text{ g/mol} = 12.816 \text{ g}
\]
\[
0.6 \text{ mol water } \times 18.016 \text{ g/mol} = 10.806 \text{ g}
\]
Total = 23.622 g \( \rightarrow x = 0.5425 \text{ wt frac methanol}
\]

Vapor: 0.729 mol methanol = 23.357 g
\[
0.271 \text{ mol water } = 4.881 \text{ g}
\]
28.238 g \( \rightarrow y = 0.8271 \text{ wt frac methanol.}
\]
Similar calculations for: 0.3 mole frac liquid give $x_{wt} = 0.433$ and $y_{wt} = 0.7793$, $T = 78.0 \, ^\circ C$

0.2 mole frac liquid give $x_{wt} = 0.3078$ and $y_{wt} = 0.7099$, $T = 81.7 \, ^\circ C$

0.15 mole frac liquid give $x_{wt} = 0.2389$ and $y_{wt} = 0.6557$, $T = 84.4 \, ^\circ C$.

Plot this data on $y_{wt}$ vs $x_{wt}$ diagram. Operating line is $y = -(L/V)x + (F/V)z$ in weight units.
Slope = - 1.857, $y = x = z = 0.45$, and $y$ intercept = $z/(V/F) = 1.286$ all in weight units.

Result is $x_{M,wt} = 0.309$, $y_{M,wt} = 0.709$ (see graph). Note that plotting only the part of the graph needed to solve the problem, the scale could be increased resulting in better accuracy. By linear interpolation $T_{drum} = 81.66 \, ^\circ C$.

---

**2.F1**

| $x_B$ | 0   | 0.1 | 0.2 | 0.3 | 0.4 | 0.5 | 0.6 | 0.7 | 0.8 | 0.9 | 1
|-------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----
| $y_B$ | 0   | .22 | .38 | .52 | .62 | .71 | .79 | .85 | .91 | .96 | 1

Benzene-toluene equilibrium is plotted in Figure 13-8 of *Perry’s Chemical Engineers Handbook*, 6th ed.

**2.F2.**

Stage 1) \( z_{f_1} = 0.4 \quad f = 1/3 \quad \text{Slope} = \frac{2/3}{1/3} = -2 \),
\[
\text{Intercept} = \frac{4}{1/3} = 1.2 \quad y_1 = 0.872 \quad x_1 = 0.164 = z_2
\]

Stage 2) \( z_{f_2} = 0.164 \quad f = 2/3 \quad \text{Slope} = -\frac{1/3}{2/3} = -1/2 \),
\[
\text{Intercept} = \frac{0.164}{2/3} = 0.246 \quad x_2 = 0.01 \quad y_2 = 0.240 = -z_3
\]

Stage 3) \( z_{f_3} = 0.240 \quad f = 1/2 \quad \text{Slope} = -1 \),
\[
\text{Intercept} = \frac{0.240}{1/2} = 0.480 \quad x_3 = 0.022 \quad y_3 = 0.461
\]

2.F3. \textit{Bubble Pt.} At P = 250 kPa. Want \( \sum K_i z_i = 1 \). Solution uses DePriester chart for K values.

Guess \( T = -18^\circ C, \quad K_1 = 1, \quad K_2 = 0.043, \quad K_3 = 0.0095, \quad \sum = 0.52 \)
Converge to \( T = 0^\circ C \)

\textit{Dew Pt. Calc.} Want \( \sum \frac{z_i}{K_i} = 1.0 \)

Try \( T = 0^\circ C, \quad K_1 = 1.93, \quad K_2 = 0.11, \quad K_3 = 0.0033, \quad \sum = 120.26 \)
Converge to \( T = 124^\circ C \). This is a wide boiling feed.

\( T_{\text{drum}} \) must be lower than 95°C since that is the feed temperature.
First Trial: Guess $T_{d1} = 70^\circ C$: $K_1 = 7.8$, $K_2 = 1.07$, $K_3 = 0.083$

Guess $V/F = 0.5$. Rachford Rice Eq.

$$f \frac{V}{F} = \frac{(7.8-1)(0.517)}{1+(6.8)(0.5)} + \frac{(0.07)(0.091)}{1+(0.07)(0.5)} + \frac{(0.083-1)(0.392)}{1+(0.083-1)(0.5)} = 0.14$$

$V/F = 0.5$ gives $f(0.5) = -0.101$

By linear interpolation: $V/F = 0.56$, $f(0.56) = -0.0016$ which is close enough for first trial.

$$V = (V/F)F = 56, \quad L = 44$$

$$x_i = \frac{z_i}{1+(K_i-1)V/F} \quad \text{and} \quad y_i = K_i x_i$$

$x_1 = 0.1075, \quad x_2 = 0.088, \quad x_3 = 0.086 \quad \sum x = 1.001$

$y_1 = 0.839, \quad y_2 = 0.094, \quad y_3 = 0.067 \quad \sum y = 0.9999$

Data: Pick $T_{ref} = 25^\circ C$. (Perry’s 6th ed; p. 3-127), and (Perry’s 6th ed; p. 3-138)

$$\lambda_1 = 81.76 \text{ cal/g } \times 44 = 3597.44 \text{ kcal/kmol}$$

$$\lambda_2 = 87.54 \text{ cal/g } \times 72 = 6302.88 \text{ kcal/kmol}$$

$$\lambda_3 = 86.80 \text{ cal/g } \times 114 = 9895.2 \text{ kcal/kmol}$$

at $T = 0^\circ C$, $C_{p1} = 0.576 \text{ cal/(g }^\circ C) \times 44 = 25.34 \text{ kcal/(kmol }^\circ C)$.

For $T = 20$ to $123^\circ C$, $C_{p1.2} = 65.89 \text{ kcal/(kmol }^\circ C)$

at $T = 75^\circ C$, $C_{p1.3} = 39.66 \text{ kcal/(kmol }^\circ C)$. (Himmelblau/Appendix E-7)

$$C_{pv} = a + bT + cT^2$$

propane $a = 16.26, \quad b = 5.398 \times 10^{-2}, \quad c = -3.134 \times 10^{-5}$

n-pentane $a = 27.45, \quad b = 8.148 \times 10^{-2}, \quad c = -4.538 \times 10^{-5}$

**n-octane $a = 8.163, \quad b = 140.217 \times 10^{-3}, \quad c = -44.127 \times 10^{-6}$

** Smith & Van Ness p. 106

Energy Balance: $E(T_d) = VH_v + LH_L - FH_F = 0$

$$F H_F = 100 \left[ (0.577)(25.34) + (0.091)(39.66) + 0.392(65.89) \right] (95.25) = 297,773 \text{ kcal/h}$$

$$L H_L = 44 \left[ (0.1075)(25.34) + (0.088)(39.66) + 0.086(16.26) \right] (70.25) = 117,450$$

$$V H_v = 56 \left[ (0.839)(3597.4 + 16.26 + 5.398 \times 10^{-2}) (45) \right] + (0.94) \left[ 6302.88 + 27.45 + 8.148 \times 10^{-2} (45) \right]

+ (0.67) \left[ 9895.3 + 8.163 \times 140.217 \times 10^{-3} (45) \right] = 240,423$$

$$E(T_{drum}) = -60,101 \quad \text{Thus, } T_{drum} \text{ is too high.}$$

Converge on $T_{drum} = 57.2^\circ C$: $K_1 = 6.4, \quad K_2 = 0.8, \quad K_3 = 0.054$

For $V/F = 0.513$, $f(0.513) = -0.0027$. $V = 51.3, \quad L = 48.7$

$x_1 = 0.137, \quad x_2 = 0.101, \quad x_3 = 0.762, \quad \sum x_i = 1.0000$

$y_1 = 0.878, \quad y_2 = 0.081, \quad y_3 = 0.041, \quad \sum y_i = 1.0000$
Thus \( T_{\text{drum}} \) must be very close to 57.3°C.

\[
\begin{align*}
    x_1 &= 0.136, \\
    x_2 &= 0.101, \\
    x_3 &= 0.762, \\
    y_1 &= 0.328, \\
    y_2 &= 0.081, \\
    y_3 &= 0.041
\end{align*}
\]

\[
V = 51.3 \text{ kmol/h}, \quad L = 48.7 \text{ kmol/h}
\]

Note: With different data \( T_{\text{drum}} \) may vary significantly.

2.F4. New Problem 4th edition. This is a mass and energy balance problem disguised as a flash distillation problem. Data is readily available in steam tables. At 5000 kPa and 500K the feed is a liquid, \( h_F = 17.604 \text{ kJ/mol} \). For an adiabatic flash, \( h_F = \left[ \frac{VH_V + Lh_L}{F} \right] \).

Vapor and liquid are in equilibrium. Saturated steam at 100 kPa is at \( T = 372.76 \text{K} \), \( h_L = 7.5214 \text{ kJ/mol} \), \( H_V = 48.19 \text{ kJ/mol} \).

Mass balance: \( F = V + L \) where \( F \) in kmol/min = (1500 kg/min)/(18.016 kg) = 83.259 kmol/min

Energy balance: \( Fh_F = VH_V + Lh_L \)

\[
\left(83.259 \text{ kmol/min}\right)(17.604 \text{ kJ/mol})(1000 \text{ mol/kmol}) = (48.19)(1000)V + (7.5214)(1000)L.
\]

Solve equations simultaneously. \( L = 62.617 \text{ kmol/min} = 1128.12 \text{ kg/min} \) and \( V = 20.642 \text{ kmol/min} = 371.88 \text{ kg/min} \)


\[
T_{\text{drum}} = 33.13 \text{ ℃}, \quad L = 34.82 \text{ and } V = 65.18 \text{ kmol/h}
\]

In order ethylene, ethane, propane, propylene, n-butane, \( x_i \) (\( y_i \)) are:

\[
(0.0122 \ (0.0748), \ 0.0866 \ (0.3005), \ 0.3318 \ (0.3781), \ 0.0306 \ (0.0404), \ 0.5388 \ (0.2062))
\]

2.G2. New problem in 4th edition. Part a. \( p = 31.26 \text{ kPa} \) with \( V/F)_{\text{feed}} = 0.0009903 \).

Part b. Use \( p_{\text{feed}} = 31.76 \text{ kPa} \), \( V/F)_{\text{feed}} = 0.018605 \),

Part c. Drum \( p = 3.9 \text{ bar} \), \( T_{\text{drum}} = 19.339 \), \( V/F) = 0.18605 \),

Liquid mole fractions: \( C_1 = 0.14663, \ C_2 = 0.027869 (\sum = 0.05253 \text{ is in spec}) \), \( C_5 = 0.6171, \ C_6 = 0.3404 \).

Vapor mole fractions: \( C_1 = 0.68836, \ C_2 = 0.20057, \ C_5 = 0.9523 \), and \( C_6 = 0.01584 \).

2.G3. New problem 4th edition. K values in Aspen Plus are higher by 17.6% (methane), 7.04% (n-butane) and 0.07% n-pentane. Since the K values are higher \( V/F \) is higher by 10.2%.

Results:

<table>
<thead>
<tr>
<th>Comp</th>
<th>x</th>
<th>y</th>
<th>K</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methane</td>
<td>0.004599</td>
<td>0.27039</td>
<td>58.79</td>
</tr>
<tr>
<td>n-butane</td>
<td>0.44567</td>
<td>0.52474</td>
<td>1.1774</td>
</tr>
<tr>
<td>n-pentane</td>
<td>0.54973</td>
<td>0.20488</td>
<td>0.37269</td>
</tr>
</tbody>
</table>

\( V/F)_{\text{drum}} = 0.43419; \ V/F)_{\text{feed}} = 0.3654; \ Q = -3183.4 \text{ cal/s} \)


<table>
<thead>
<tr>
<th>COMP</th>
<th>x(l)</th>
<th>y(l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>METHANE</td>
<td>0.12053E-01</td>
<td>0.84824</td>
</tr>
<tr>
<td>BUTANE</td>
<td>0.12978</td>
<td>0.78744E-01</td>
</tr>
<tr>
<td>PENTANE</td>
<td>0.29304</td>
<td>0.47918E-01</td>
</tr>
<tr>
<td>HEXANE</td>
<td>0.56513</td>
<td>0.25101E-01</td>
</tr>
<tr>
<td>V/F = 0.58354</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

2.G5. New. Used NRTL. \( T = 368.07, \ Q = 14889 \text{ kW}, 1^{st} \text{ liquid/total liquid} = 0.4221,\)

<table>
<thead>
<tr>
<th>Comp</th>
<th>Liquid 1, x₁</th>
<th>Liquid 2, x₂</th>
<th>Vapor, y</th>
</tr>
</thead>
</table>

51
2.G6. Used Peng Robinson. Feed pressure = 10.6216 atm, Feed temperature = 81.14°C, V/F = 0.40001, Q_drum = 0. There are very small differences in feed temperature with different versions of AspenPlus.

<table>
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<tr>
<th>COMP</th>
<th>x(l)</th>
<th>y(l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>METHANE</td>
<td>0.000273</td>
<td>0.04959</td>
</tr>
<tr>
<td>BUTANE</td>
<td>0.18015</td>
<td>0.47976</td>
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<tr>
<td>PENTANE</td>
<td>0.51681</td>
<td>0.39979</td>
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<tr>
<td>HEXANE</td>
<td>0.30276</td>
<td>0.07086</td>
</tr>
<tr>
<td>V/F = 0.40001</td>
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<td></td>
</tr>
</tbody>
</table>


2.H3. New Problem. 4th ed. Answer V/F = 0.564; xE = 0.00853, xhex = 0.421, x hept = .570; yE = .421, y Hex = 0.378, y Hept = .201.

2H4. New Problem, 4th ed. Answer: p_drum = 120.01, kPa = 17.40 psia
 xB = 0.1561, xpen = 0.4255, x hept = 0.4184, yB = 0.5130, yPen = 0.4326, yhept = 0.0544

 a. SOLUTION. P = 198.52 kPa.
 b. V/F = 0.24836, ethane x = 0.00337, y = 0.0824; Propane x = 0.05069, y = 0.3539;
 Butane x = 0.1945, y = 0.3536; Pentane x = 0.3295, y = 0.1584; Hexane x = 0.3198, y = 0.0469
 Heptane x = 0.1022, y = 0.00464
 c. T = 34.48°C
 d. T = -1.586°C and V/F = 0.0567


<table>
<thead>
<tr>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
<th>E</th>
<th>F</th>
<th>G</th>
<th>H</th>
<th>I</th>
<th>J</th>
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</tr>
<tr>
<td>2</td>
<td>Example 2-2 on spreadsheet</td>
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<td>K4</td>
<td>0.3</td>
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<td>z3</td>
<td>0.15</td>
<td>z4</td>
<td>0.45</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>Guess V/F</td>
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<td></td>
<td></td>
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<td></td>
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<td></td>
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<tr>
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<td>x2</td>
<td>0.058784</td>
<td>x3</td>
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<td>x4</td>
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</tr>
<tr>
<td>7</td>
<td>y1</td>
<td>0.524353</td>
<td>y2</td>
<td>0.141081</td>
<td>y3</td>
<td>0.12</td>
<td>y4</td>
<td>0.207877</td>
<td>sum ↓</td>
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<td>-0.48505</td>
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<tr>
<td>9</td>
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<td>Goal seek 9! to zero by changing B5</td>
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<td>-0.00021</td>
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chk