Solution Manual for Separation Process Engineering Includes Mass Transfer Analysis 4th Edition Wankat 0133443655 9780133443653 Link full download Solution Manual:

https://testbankpack.com/p/solution-manual-for-separationprocess-engineering-includes-mass-transfer-analysis-4thedition-wankat-0133443655-9780133443653/ *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.

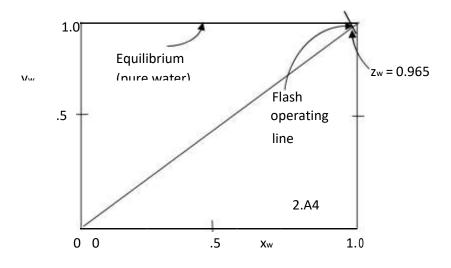
2.A1. Feed to flash drum is a liquid at high pressure. At this pressure its enthalpy can be calculated as a liquid. eg. h $T_{F,Phigh}$ $c_{p \ LIQ}$ T_{F} T_{ref} . When pressure is dropped the mixture is above

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.A2. Yes.

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

2.A4.



2.A6. In a flash drum separating a multicomponent mixture, raising the pressure will:i. Decrease the drum diameter and decrease the relative volatilities. *Answer is i.*

2.A8. New Problem in 4th ed.

. 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

c. Within a homologous series such as light hydrocarbons as the molecular weight increases, the K value (at constant pressure and temperature)

b

- a. increases, b. decreases, c. stays constant b
- d. At what pressure does pure propane boil at a temperature of -30_{\circ} C? 160 kPa

- 2.A9. a. The answer is <u>3.5 to 3.6</u> b. The answer is <u> $36^{\circ}C$ </u> c. *This part is new in 4th ed.* <u> $102^{\circ}C$ </u>
- 2.A10. Parts c, d, and e are new in 4^{th} 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* 4^{th} *edition.* Although V is an extensive variable, V/F is an intensive variable and thus satisfies Gibbs phase rule.
 - 2A14. New problem in 4^{th} 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:

$\mathbf{F}, \mathbf{Z}, \mathbf{T}, \mathbf{P}_{\text{drum}}, \mathbf{P}_{\text{drum}}$	F,T _F ,z,p	F, h _F , z,p
F,z, y, P _{drum}	F,T _F ,z, y	F, h _F , z, y
F,z, x,p _{drum}	F,T _F ,z, x	etc.
F,z, y,p _{drum}	F, T_F, z, T_{drum} , p_{drum}	
F,z, x,T _{drum}	F,T _F , y,p	
Drum dimensions, z , F_{drum} , p_{drum}	F,T _F , y,T _{drum}	
Drum dimensions, z, y,p _{drum}	F,T _F , x,p	
etc.	F,T_F , x,T_{drum}	
	F,T _F , y, x	

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 $\frac{\text{lb mole}}{\text{hr}}$, D .98 and L 2.95 ft from Problem 2-D1e.

Since D $\alpha \sqrt{V}$ and for constant V/F, V α F, we have D $\alpha \sqrt{F}$. With F = 25,000:

$$\sqrt{F_{\text{new}}}/\overline{F_{\text{old}}} = 5$$
, $D_{\text{new}} = 5 D_{\text{old}} = 4.90$, and $L_{\text{new}} = 3 D_{\text{new}} = 14.7$.

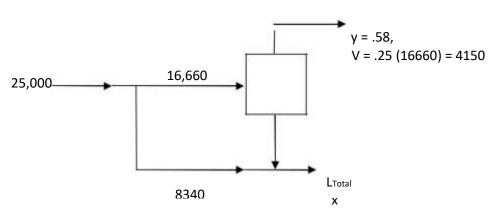
Existing drum is too small.



Alternatives

a) Do drums in parallel. Add a second drum which can handle remaining 8340 lbmol/h.

b) Bypass with liquid mixing



Since x is not specified, use bypass. This produces less vapor.

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

$$D \sqrt{\frac{V MW_{v}}{3K_{drum} 3600 L v v}}$$

Bypass reduces V

- c1) K_{drum} is already 0.35. Perhaps small improvements can be made with a better demister \rightarrow Talk to the manufacturers.
- c2) ρ_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.

V

2.C2.

$$\begin{array}{cccc} F & K_B & 1 & K_A & 1 \\ x & \underline{Fz_i} & & \end{array}$$

2.C5. a. Start with $$^{\rm i}$$ $$L_{\rm VK_i}$$ and let V F L

$$x_{i} \quad \frac{Fz_{i}}{L F L K_{i}} \quad \text{or } x_{i} \frac{Z_{i}}{L} \quad L_{\kappa_{i}}$$

$$F \quad F$$

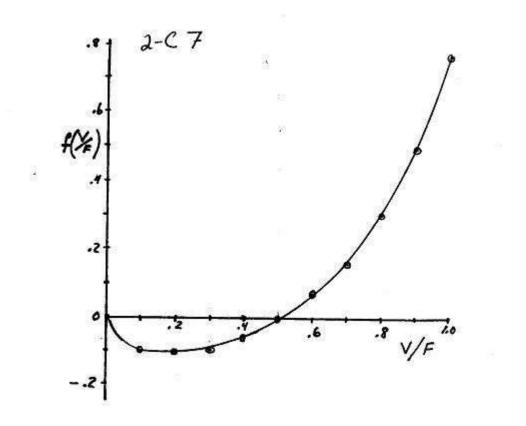
Then $y_i K_i x_i \frac{K_i z_i}{\frac{L}{F} - 1} \frac{K_i}{F}$ From $y_i x_i = 0$ we obtain $\frac{\frac{K_i}{L} \frac{1}{F} \frac{z_i}{F}}{F}$

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.

F

SOLUTION:
y
$$\frac{L}{V}$$
 y $\frac{F}{V}$ z.rearrange:
y $\frac{L}{V}$ $\frac{F}{V}$ z.rearrange:
therefore
x = y = z (2-18)

The intersection is at the feed composition.



2.C8. Derivation of Eqs. (2-62) and (2-63). Overall and component mass balances are,

 $\begin{array}{ccccccc} F \ V \ L_1 & L_2 \ \text{and} \ Fz \ i \ L_1 \ x \ _{i,L1} & L_2 & x \ _{i,L2} Vy_i \ \ \text{Substituting in Eqs. (2-60b) and 2-60c)} \\ & Fz \ L \ K & x & L \ x & VK \ x \end{array}$

i 1 i,L1 L2 i,L 2 2 i,L2 iV L 2 i,L 2

Solving,

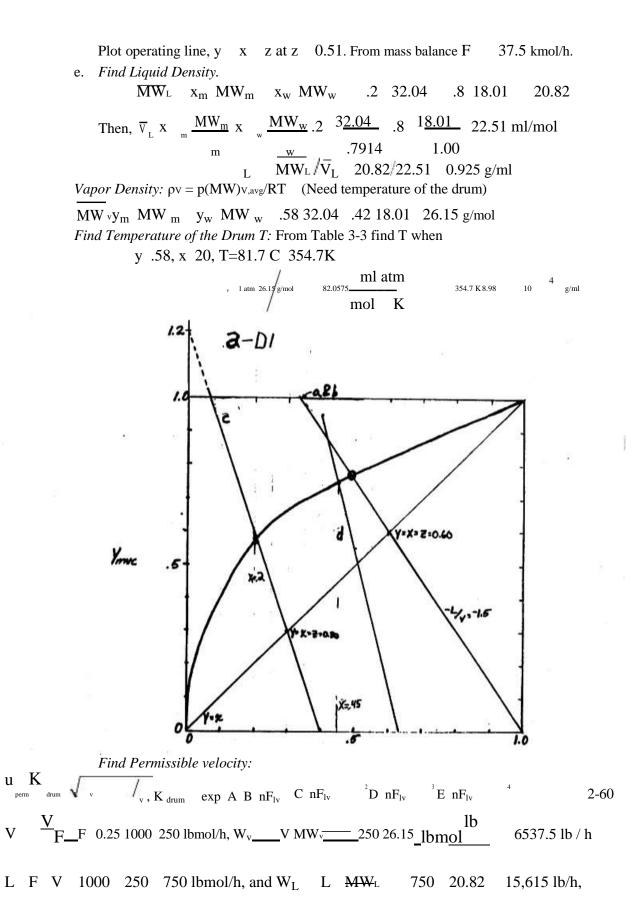
Dividing numerator and denominator by F and collecting terms.

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_{int ercept} 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
$$\frac{L}{V} = \frac{F V}{V} = \frac{1 V F}{V/F} = \frac{.8}{.2} 4$$



$$F_{1v} = \frac{W}{W_v} \sqrt{\frac{1}{L}} = \frac{15615}{6537.5} = \frac{8.89}{.925} \frac{10^4}{9.25} = 0.0744, \text{ and n } F_{1v}2.598$$
Then K dammed and the problem is the second structure of the system o

x ethane	T °C	y ethane
0	63.19	0
.025	56.18	0.1610
.05	49.57	0.2970
.10	37.57	0.5060
.15	27.17	0.6503
.20	18.26	0.7492
.25	10.64	0.8175
.30	4.11	0.8652
1.0	-37.47	1.0

2.D3. New Problem in 4^{th} edition.. Part a.

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 $T = 18.26 \,^{\circ}$ 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 \,^{\circ}C (highest).

c. See figure. Slope = -L/V = -(1 - V/F)/(V/F) = -.6/.4 = -1.5. xE = 0.12, yE = 0.57, T = 33.4° C.d. From equilibrium data yE = 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)(0.1 - 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 3^{rd} edition. Highest temperature is dew point V **F** 0

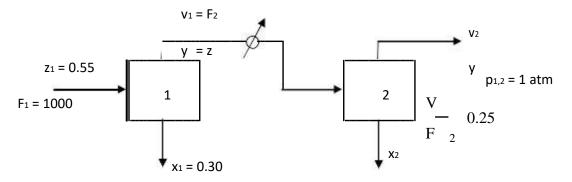
Set

If pick C4 as reference: First guess butane 1.0, T 41 C : K_{C3} 3.1,K_{C6} 0.125

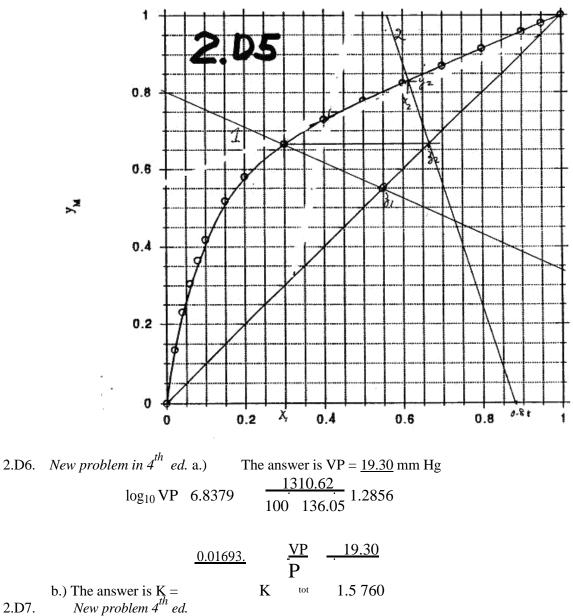
$$\begin{array}{c} \underbrace{y \ .2 \ .35 \ .45}_{-i} & 1.20 \ \mathrm{K_{i}} \ 6 \\ 2.45 \ .44 \\ & \mathrm{K_{C4,NEW}} \ 2.45 \ 1.2 \\ & 2.94, \ T \ 96 \ \mathrm{C} : \mathrm{K_{C3}} \ 6.9, \ \mathrm{K_{C6}} \ 0.56 \\ & \underbrace{y \ .2 \ .35 \ .45}_{-i} & 0.804 \ \mathrm{Gives} \ 84 \ \mathrm{C} \ \mathrm{K_{i}} \ 6.9 \\ & \underbrace{\mathrm{Use} \ 90.5^\circ \rightarrow \mathrm{Avg} \ \mathrm{last two} \ \mathrm{T} \ \ \mathrm{K_{C4}} \ 2.7, \ \mathrm{K_{C3}} \ 6.5, \ \mathrm{K_{C6}} \ 0.49 \\ & \mathrm{y_i} \ / \mathrm{K_i} \ \ 6.5 & \underbrace{.2 \ .35}_{-2.7} \ .49 & 1.079, \ \mathrm{T} \sim 87 \ 88^\circ \ \mathrm{C} \end{array}$$

Note: hexane probably better choice as reference. a)

2.D5.



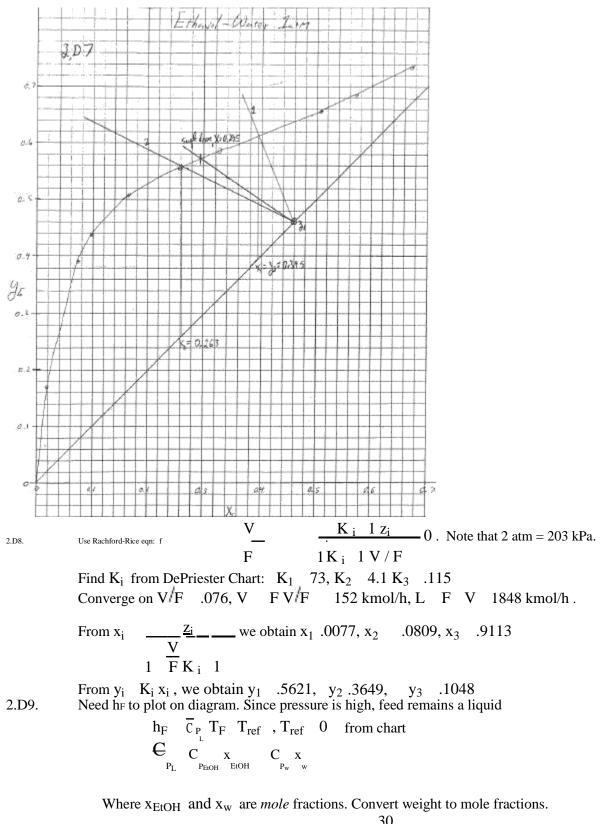
b)
$$\frac{y_{1V}}{y_{1V}} = \frac{F}{x_{1} + v_{2} + Plot 1 \text{ Op line.}}$$
 st $y_{1} = 0.66 = z_{2}$
 $y = x = z = 0.55 \text{ to } x_{1} = 0.3 \text{ on eq. curve (see graph)}$
Slope $\frac{L}{0.55} = 0.25 + 0.25 + 0.454545 \text{ L} = V + F + 1000}{V_{1}} = \frac{V - 687.5}{.55} = 0.454545 \text{ L} = V + F + 1000}{V_{1}} = \frac{V - 687.5}{.55} = 0.6875}{V_{1} = 687.5 \text{ kmol/h} = F_{2}}$
c) Stage 2 $\frac{V}{.} = 0.25$, $\frac{L}{0.75F} = 0.75F}{V_{1} = 0.075F} = \frac{V - 687.5}{.5} = 0.666}{V_{1} = 0.25F}$
At x 0, y z $\frac{V}{V} = \frac{0.66}{0.25} = 2.64$. At y 0, $x_{2} = \frac{F}{L} = \frac{Z}{L'} = \frac{0.66}{0.75} = 0.88$
From graph $\frac{V - 0.82}{.5} = \frac{V - 0.25}{.5} = \frac{V - 0.25}{.5} = \frac{171.875 \text{ kmol/h}}{.55} = \frac{F_{2}}{.55} = \frac{V - 0.25}{.55} = \frac{171.875 \text{ kmol/h}}{.55} = \frac$



2.D7.

Part a. Drum 1: $V_1/F_1 = 0.3$, Slope op line = -L/V = -.7/.3 = -7/3, $y=x=z_1=0.46$. $L_1 = F_2 = -.7/3$ 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.

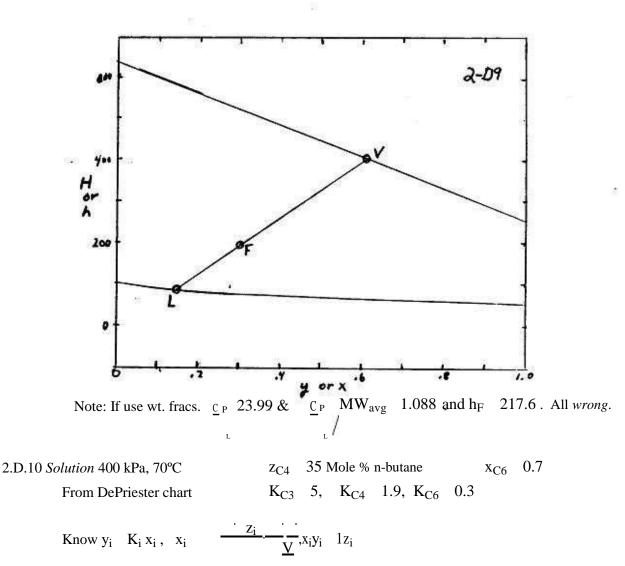


Basis: 100 kg mixture: 30 kg EtOH $\frac{30}{46.07}$ 0.651 kmol

70 kg water 70/18.016 3.885 Total = 4.536 kmol

Avg. MW $\frac{100}{4.536}$ 22.046 Mole fracs: $x_E = \frac{0.6512}{4.536}$ 0.1435, $x_w = 0.8565$. Use $C_{P_{L_{EOH}}}$ at 100 C as an average C_P value. $\overline{C}_P = 37.96$.1435 18.0 .8565 20.86 kcal L kmol C Per kg this is $\frac{\underline{C}}{\underline{P}_L} = \frac{20.86}{20.046} = 0.946$ kg C $h_F = 0.946$ 2000 189.2 kcal/kg

which can now be plotted on the enthalpy composition diagram. Obtain T_{drum} 88.2 C, x_E 0.146, and 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



– 1 K_i 1 F R.R. $\frac{K_{i} \ 1 \ z_{i}}{\sum_{C_{6}}^{V} K_{i} \ 1 \ F} = 0 \quad z_{C3} \ 1 \ z_{C6} \ z_{C4} \ .65 \ z_{C6}$

C6: 0.7
$$\underline{V} = \underline{V}^{z_{C6}} = 0.71 \ 0.7 F^{z_{C6}} = 0.7 \ 0.49 F^{z_{C6}}$$

 $1 \ K_{C6} \ 1 \ F \ 1 \ 0.7 \ F$

RR Eq:
$$\frac{4.65 \text{ z}_{C6}}{1.4 \text{ }\frac{\text{V}}{\text{F}}} = \frac{0.9.35}{1.0.9 \text{ }\frac{\text{V}}{\text{F}}} = \frac{0.7 \text{ z}_{C6}}{10.7 \text{ }\frac{\text{V}}{\text{F}}} = 0$$

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

2.D11. New Problem 4^{th} ed. Obtain K ethylene = 2.2, K propylene = 0.56 from De Priester chart.

$$\begin{split} &K_E = y_E/x_E \text{ and } K_P = y_P/x_P \text{ Since } y_P = 1 - y_E \text{ and } x_P = 1 - x_E \text{ , } K_P = (1 - y_E)/(1 - x_E). \\ &Thus, 2 \text{ eqs and } 2 \text{ unknowns. Solve for } y_E \text{ and } x_E. \\ &x_E = (1 - K_P) / (K_E - K_P) \text{ and } y_E = K_E \text{ } x_E = K_E (1 - K_P) / (K_E - K_P) \\ &x_E = (1 - 0.56) / (2.2 - 0.56) = 0.268 \text{ and } y_E = K_E \text{ } x_E = (2.2)(0.268) = 0.590 \\ &Check: x_P = 1 - x_E = 1 - 0.268 = 0.732 \text{ and } y_P = 1 - y_E = 1 - 0.590 = 0.410 \\ &K_P = y_P/x_P = 0.410 / 0.732 = 0.56 \text{ OK} \end{split}$$

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. \overline{MW}_{L} x_{m} MW_{m} x_{w} MW_{w} .2 32.04 .8 18.01 20.82 $\overline{v}_{L} x_{m} \frac{MW_{m}}{m} x_{w} \frac{MW_{w}}{m} .2 \frac{32.04}{.8} .8 \frac{18.01}{.100} 22.51 \text{ ml/mol}$ $L MW_{L} v_{L} 20.82 22.51 0.925 \text{ g/ml}$ Then, $\frac{p MW}{RT}$ (Need temperature of the drum) Find Vapor Density. v $MW \ _v y_m \ MW \ _m \ y_w \ MW \ _w \ .58 \ 32.04 \ .42 \ 18.01 \ \ 26.15 \ \text{g/mol}$ Find Temperature of the Drum T: 2-7 find Т corresponding From Table to y .58, x 20, T=81.7 C 354.7K / 1 atm 26.15 g/mol 82.0575 ml atm mol K 8.98 10 4 g/ml

Find Permissible velocity: $u_{perm} = K_{drum} = L_{v} = v$

L F V 1000 250 750 lbmol/h, and W_L L MW_L 750 20.82 15,615 lb/h,

With L/D = 4, D
$$\sqrt{4A_T 3.41}$$
 ft and L 13.6 ft

2.D13. New Problem 4^{th} ed. Xbutane = $1 - x_E = 0.912$, ybutane = $1 - y_E = 0.454$. Ke = yE/xE = 0.546/0.088 = 0.546/0.088

6.20, K_{butane} = $y_B/x_B = 0.454/0.912 = 0.498$.

Plot K_E and K_{butane} on DePriester chart. Draw straight line between them. Intersections with T and P axis give $T_{drum} = 15$ °C, and $p_{drum} = 385$ 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. New Problem 4^{th} ed. DePriester chart, Fig. 2-12: K_{C1} = 50, K_{C4} = 1.1, and K_{C5} = 0.37; z₁ = 0.12, z₄ = 0.48, z₅ = 0.40

Rachford-Rice equation:
$$\frac{K_{C2}}{V} \frac{1}{2} \frac{z_{C1}}{V} \cdot \frac{K_{iC4}}{V} \frac{1}{z_{nC4}} \frac{K_{nC4}}{V} \frac{1}{z_{nC5}} \frac{0}{0} \frac{V}{V}$$

$$1 K_{C1} 1 F 1 K_{nC4} 1 F 1 K_{nC5} 1 F$$

$$5.88 - 0.048 - 0.252$$

Equation becomes: $1 \frac{49(V/F)}{1 0.1(V/F)} = 1 \frac{0.63(V/F)}{0} = 0$

Trials: V/F = 0.4, Eq. = -.005345; V/F = 0.39, Eq. = 0.004506; V/F = 0.394, Eq. = 0.000546, which is close enough with DePriester chart.

Liquid mole fractions:

X c1 .12
$$x_{c4} = 0.4618, x_{c5} = 0.5321, and $\sum x_i = 0.9998$$$

 $^{C1} 1 K_{C1} 1 (V / F) 1 49.394 0.00591;$

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: zc2, zic4, znc4. Need two other
 - eqns: z_{iC4}/z_{nC4} constant, and z_{C2} z_{iC4} z_{nC4} 1.0 Thus, solve three equations and three unknowns simultaneously. *Do It.* Rachford-Rice equation is,

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.

$$Z_{nC4} - \underbrace{\frac{K_{C2} 1}{1 K_{C2} 1 F}}_{1.8 K_{C2} 1 F} - \underbrace{\frac{K_{nC4} 1}{1 K_{C2} 1 F}}_{1 K_{C2} 1 F 1 K_{iC4} 1 F} \underbrace{\frac{K_{nC4} 1}{V}}_{1 K_{nC4} 1 F}$$

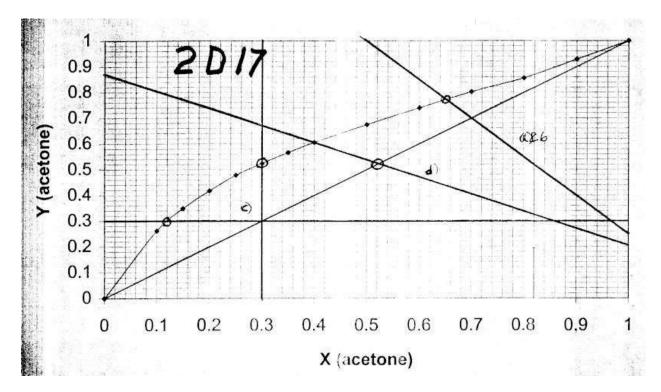
Thus,

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 1^{st} guess. Can assume all C₁ in vapor, ~ 1/3 C₄ in vapor, C₅ & C₆ in bottom

Similar for other components.

- 2-D17. a. V 0.4F 400, L 600 SlopeL/F 1.5
 - Intercepts y = x = z = 0.70. Plot line and find $x_A = 0.65$, $y_A = 0.77$ (see graph) b. V = 2000, 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, L = 400, -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
$$Z_{\underline{h}} = 1$$

 $i \qquad \underline{Z_{\underline{h}}} = 1$
 $V \qquad \underline{X_{\underline{h}}} = 1$

Guess T_{drum} , calculate K_h , K_b and K_p , and then determine V/F.

Check:
$$\frac{K_1}{1 K_1} \frac{1 z_i}{1 V F} 0?$$

Initial guess: T_{drum} must be less than temperature to boil pure hexane

 K_h 1.0, T 94 C . Try 85°C as first guess (this is not very will tell critical and the calculation us if there is a mistake). K_h =0.8, K_b 4.8, K_p =11.7.

$$\frac{0.6}{V} = \frac{0.6}{0.851} = 1.471. \text{ Not possible. Must have } K_h = 0.85 = 0.706$$
F = 0.8 1
Try T 73 C where $K_h = 0.6. \text{ Then } K_b = 3.8, K_p = 9.9.$

$$\frac{V}{F} = \frac{0.6}{.851} = 0.735$$
F = .6 1

Check:

Solve for

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}

equations and two unknowns. Remove x_{C6} and solve

$$z_{C6} \cdot .93C_{6} - \frac{.9 z_{C6} \text{ KV}}{1 \text{ V/F}} \text{ F}$$

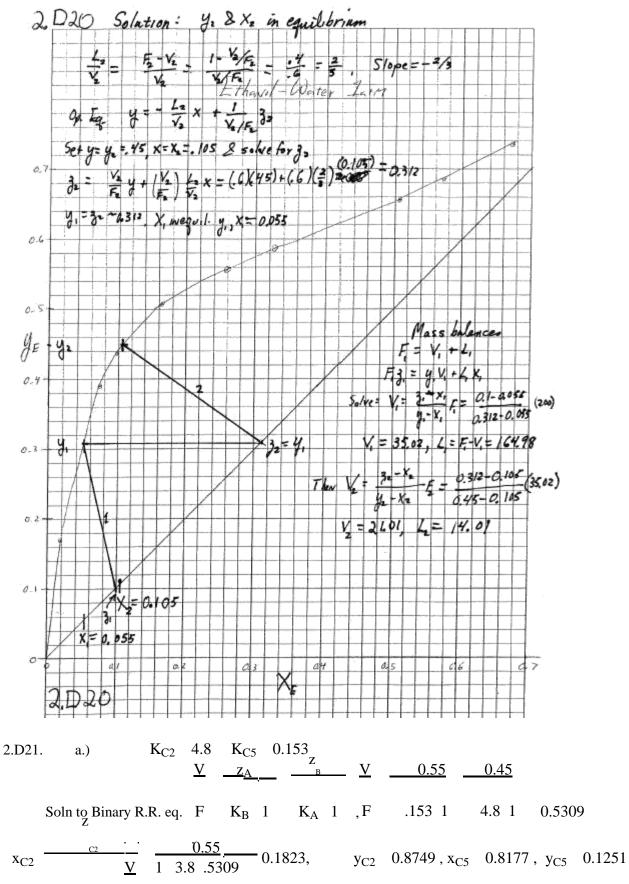
$$\frac{\text{V}}{\text{F}} \cdot .1$$

$$\text{V F.} \quad \text{F} \cdot .9K_{C6} \quad .1 \quad . \text{ Trial and error scheme.}$$

Pick T, Calc K_{CK} Calc V F, and Check f V F^l 0? If not K ref_{new} ref_{old} 1 df TTry **F** .9 .37 .1 0.231. **Rachford Rice equation** 2.1.4 .08 .25 .63.35 1 .08 .231 1 2.1.231 1 .63 .231 .28719 f ΚΤ .37 ^{new} 1 0.28719 0.28745 use .28 ref

Converge on $T_{New} \sim 57$ C. Then $K_{C4} = 2.50$, $K_{C8} = .67$, and V/F = 0.293.

2.D20. New Problem 4^{th} ed.



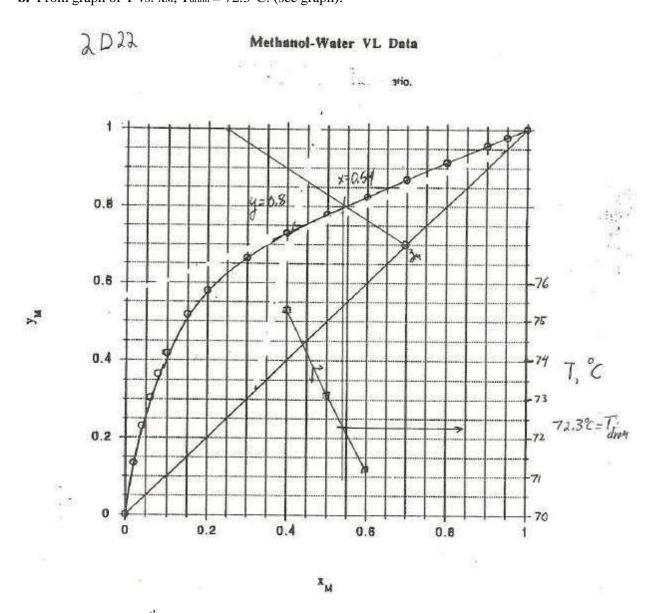
1 K_{C2} 1 F

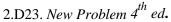
Need to convert F to kmol. Avg MW 0.55 30.07 0.45 72.15 49.17 F 100,000 $\frac{\text{kg}}{\text{hr}} \left| \frac{\text{kmol}}{49.17 \text{ kg}} \right|$ 2033.7 kmol/h , V V F/F 1079.7, L F V 954.0 kmol/h $u K_{Perm drum} \sqrt{\frac{Lv}{v}}$ b.) MW L0.1823 30.07 0.8177 72.15 64.48 To find MWv0.8749 30.07 0.1251 72.15 35.33 For liquid assume ideal mixture: $\overline{\nabla} x \quad \overline{\nabla} \quad x \quad \overline{\overline{\nabla}} \quad x \quad \overline{\underline{W}} \quad \underline{\underline{W}} \quad \underline{\underline{C}}^2 \quad x \quad \overline{\underline{MW}} \quad \underline{\underline{C}}^2$ C2 C2,liq C5 C5,liq C2 C2,liq C5 C5,ljq 1 VL 0.1823 <u>30.07</u> 0.8177 <u>72.15</u> 103.797 ml/mol 0.54 0.63 L <u>MW L 64.48</u> 0.621 g/ml V 103.797 $\frac{1}{MW_v} = \frac{101.3 \text{ kPa}}{101.3 \text{ kPa}} = \frac{1}{101.3 \text{ kPa$ For vapor: ideal gas: 82.0575 mol K 303.16K K_{drum} : Use Eq. (2-60) with $F_{IV} = \frac{W_L}{W_V} \sqrt{\frac{v}{v}}$ $W_{L} 997.7 \quad \frac{\text{kmol}}{\text{h}} \left| \frac{64.48 \text{ kg}}{\text{kmol}} \right| 6,4331.7 \text{ kg/h}, W_{V} 881.5333 31,143.4 \text{ kg/h}$ $F^{\text{IV}} = \frac{64331.7}{31,143.3} + \frac{0.009814}{0.621} = 0.2597$ K_{drum} exp 1.877478 0.81458 n.2597 0.18707 n.0.2597² kmol Α С 0.009814 h cm 1000g S m D $\sqrt{4A_{C}I.33}$ m . Arbitrarily L/D 4, L 5.32 m

^{2.}D22. New problem in 4th edition.

a. V = F - L = 50 - 20 = 20 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. xM, Tdrum = 72.3°C. (see graph).





Part a. $F_{new} = (1500 \text{ kmol/h})(1.0 \text{ lbmol/}(0.45359 \text{ kmol})) = 3307 \text{ lb mol/h}$. V, Wv, L, and WL are the values in Example 2-4 divided by 0.45359. The conversion factor divides out in F_{1v} term. Thus, F_{1v}, K_{drum}, and u_{perm} are the same as in Example 2-4. The Area increases because V

increases: Area = Area_{Example 2-4}/0.45359 = 16.047/0.45359 = 35.38 ft². *Diameter* $\sqrt{4 \text{ Area }}$ 6.71 *feet* Probably round this off to 7.0 feet and use a drum height of 28 feet.

b. $F_{parallel} = 3307 - 1500 = 1807 \ lbmol/h.$

Fiv, Kdrum, and uperm are the same as in Example 2-4. $V_{parallel} = (V/F) F_{parallel} = 0.51 (1807) = 921.6 \text{ kmol/h}$.

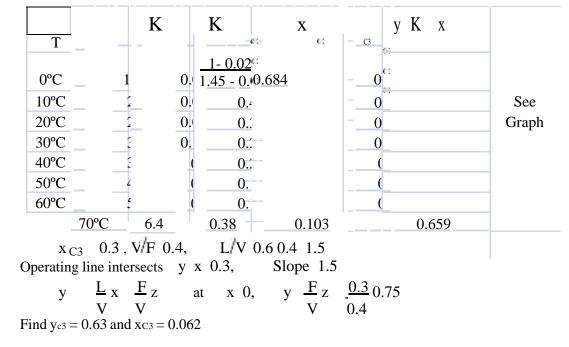
$$V = \frac{V}{16.047} = \frac{V}{V = \frac{Parallel}{Example_2 4}} = 16.047 \quad (921.6 / 765) \quad 19.33 \text{ ft}^2$$

Then, *Diameter* $\sqrt{4}$ *Area* /4.96 *feet*, Use a 5.0 feet diameter and a length of 20 feet.

2.D24. p = 300 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 1st equation into 2nd K_{C6} 1 K_{C3} x_{C3} / 1 x_{C3}
Solve for x_{C3} 1 x_{C3} K_{C6} 1 K_{C3} x_{C3} , x_{C3} K_{C6} 1 K_{C6}
 x_{C3} K_{C6}^{1} K_{C6} y_{C3} K_{C3} K_{C6} 1 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} $\frac{1 K_{C6}}{K_{C6}}$ $\frac{1 K_{C6}}{K_{C6}}$ 1.0
 $1 K_{C6}$ $\frac{K_{C6}}{K_{C6}}$ $\frac{K_{C6}}{K_{C3}}$ $\frac{K_{C3}}{K_{C3}}$ $\frac{1}{K_{C6}}$ 1.0
 $1 K_{C6}$ K_{C6}^{C3} $\frac{K_{C6}}{K_{C3}}$ $\frac{K_{C6}}{K_{C3}}$ $\frac{K_{C6}}{K_{C3}}$ $\frac{K_{C6}}{K_{C3}}$

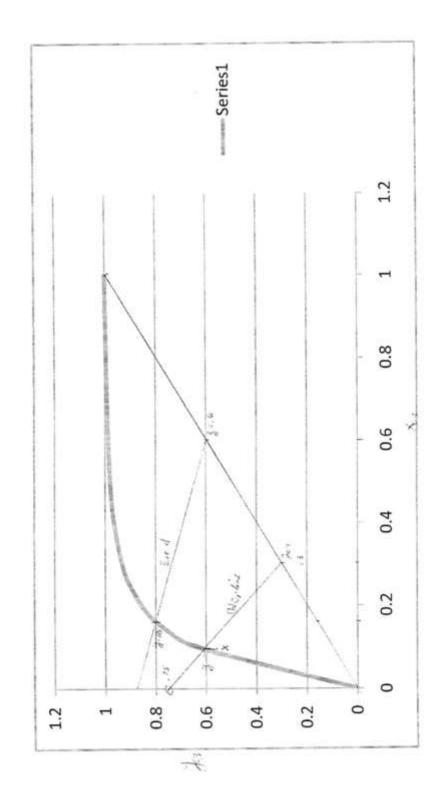
Pick intermediate temperatures, find K_{C3} $\ \&\ K_{C6}$, calculate $x_{C3}\&\ y_{C3}$.

b.



Check with operating line: 0.63 1.5 .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°C $\frac{L}{V}$ $\frac{v}{x}$ $\frac{.8.6}{.16.6}$ d. y .8, x ~ .16 Slope V/F = f = 1/1.45 = 0.69

 $0.45 \qquad \frac{1 \text{ f}}{\text{f}}.45$



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2.D25. 20% Methane and 80% n-butane. T
$$50 \, {}^{\circ}{\rm C}$$
, $\frac{V}{F}$ 0.40, Find p $_{drum}$
 $0 \, {}^{\circ}{\rm c}^{-V} - \frac{K_{\rm A}}{V} \frac{1 \, z_{\rm A}}{V} - \frac{K_{\rm B}}{F} \frac{1 \, z_{\rm B}}{V}$
F $+ K_{\rm A} + \frac{1}{V} - 1 \, K_{\rm B} + 1 \, \frac{V}{F}$
Pick pairum 1500 kPa: Kc4 13 KnC4 0.4
(Any pressure with Kc1 + 1 and Kc4 + 1.0 is 0K)
Trial 1 f $-\frac{12.2}{112.4} \frac{6.8}{1.6.4} \frac{0.2178}{0.2178}$ Need lower p $_{drum}$
K $\frac{K_{\rm C4}}{P_{\rm new}} \frac{K_{\rm C4}}{1} \frac{1}{1} \frac{6.4}{1} \frac{0.4}{1} - \frac{12.138}{2.138} 0.511$ with d = 1.0
P1160 K = 16.5, f 15.5.2 - $\frac{-489.8}{1} \frac{0.4305}{1} + \frac{4863}{2} 0.055769}$
K c4 Pnew $\frac{-0.511}{1} \frac{0.55769}{1} 0.541$, Pnew 1100, Kc1 + 17.4
f $\frac{164.2}{1} - \frac{459.8}{1} + \frac{0.0159}{2} + \frac{0.02645}{2} \frac{1}{1} \frac{16.4}{4} + \frac{459.4}{2} \frac{0.02645}{2}$
b.) $x_i - \frac{Z_i}{V} \, x_{\rm C1} - \frac{0.2}{1} \frac{0.02}{16.4} - \frac{0.02645}{2} \frac{1}{1} \frac{1}{16.4} \frac{1}{4} + \frac{459.8}{2} \frac{1}{2} \frac{0.02645}{2}$
L K₁ 1 F $\frac{1}{Y} \, y_{\rm C1} \, K_{\rm C1} \, x_{\rm C1} + \frac{1}{1} \frac{0.02}{4} \frac{1}{2} \frac{0.02645}{2} \frac{1}{2} \frac{1}{1} \frac{1}{2} \frac{$

c.) P_{tot} 191.97 mm Hg [at boiling for pure component P_{tot} VP] d.) C5: \log_{10} VP 6.853 $\frac{1064.8}{30\ 233.01}$ 2.8045 , VP 637.51 mm Hg K_{C5} VP_{C5}/P_{tot} 637.51/500 1.2750

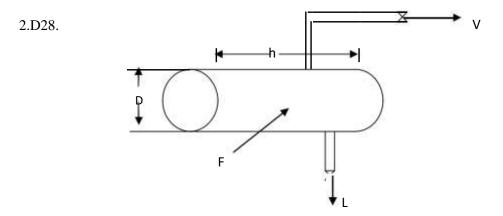
C6:
$$\log_{10}$$
 VP 6.876 $\frac{1171.17}{30}$ 2.2725 , VP 187.29 mm Hg
K_{C6} 187.29/500 0.3746
e.) K_A y_A/x_A K_B y_B/x_B (1 y_A)/(1 x_A)

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

$$X_{C5} = \frac{1}{K_{C5}} \frac{K_{C6}}{K_{C6}} = \frac{10.3746}{1.2750} \frac{10.3746}{0.3746} \frac{10.3746}{0.6946}$$

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

g.) Same as part f, except units are mol/min.

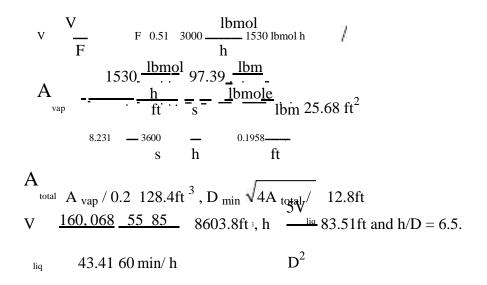


From example 2-4, $x_H = 0.19$, $T_{drum} = 378K$, V/F = 0.51, $y_H = 0.6$, $z_H = 0.40$ MW_v = 97.39 lbm/lbmole (Example 2-4)

$$\begin{array}{c|c} 3.14 & 10^{3} \text{ g mol} \\ \hline 1 \\ 454 \text{ g'lbm} \\ \hline 454 \text{ g'lbm} \\ \hline 1 \\ \text{ft}^{3} \\ \text{ft}^{3} \\ \hline 1 \\ \text{ft}^{3} \\ \text{ole 198} \\ \text{ft}^{3} \\ \hline 1 \\ \$$

$$u_{\text{perm drum }} K_{\text{drum }} \sqrt{\frac{Lv}{v}}, K_{\text{horiz}} 1.25 \text{ K}_{\text{vertical}}$$

From Example 2-4, $K_{vertical} = 0.4433$, $K_{horiz} = 1.250.4433$ 0.5541 $u_{perm} = 0.5541 + 0.6960 + 0.00314 + 12 + 0.00314 + 12 + 0.00314 + 12 + 0.00314 + 0.00$



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 $^{\circ}$ C.

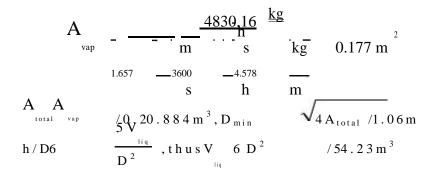
The vapor mole fractions are: methane = 0.8296, propane = 0.1458, n-butane = 0.0143, i-butane = 0.0097, and n-pentane = 0.0006.

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 = $0.004578 \text{ g/cc} = 4.578 \text{ kg/m}^3$, 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)

$$F_{1v} = (W_L/W_V) [\rho_V/\rho_L]^{0.5} = (10169.84/4830.16) [0.004578/0.60786]^{0.5} = 0.18272$$

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



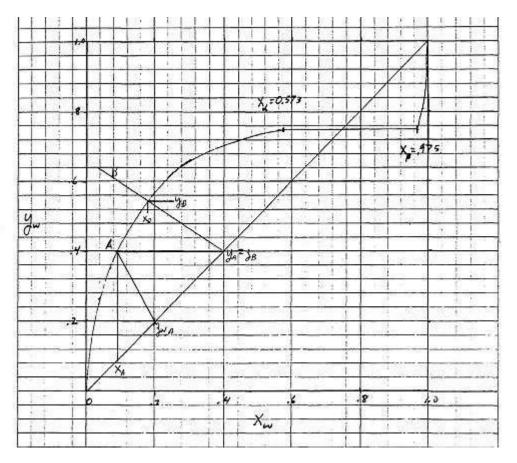
V (Volrate)(holdtime+surgetime)=(
$$\frac{10169.84 \text{ kg/h}}{607.86 \text{ kg/m}})(9/60 \text{ st})$$

st 607.86V_{lig}/10169.84 9/60 0.103 hours 6.18 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,

 $y = -(L/V)x + F_B/V)$ zB 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 $x_B = 0.18$ & $y_B = 0.54$. LB = (fraction condensed)(feed to B) = 0.4(35.48) = 14.19 kmol/h and VB = FB - LB = 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.

2.D31. New problem in 4th US edition. Was 2.D13 in 3rd International Edition.

a) Since K's are for mole fractions, need to convert feed to mole fractions. <u>Basis</u>: 100 kg feed

$$50 \text{ kg } \underline{\text{n } \text{C}_{4}} \frac{1 \text{ kmol}}{58.12 \text{ kg}} = 0.8603 \text{ kmol} \qquad | z = 0.555 \text{ z}^{4}$$

$$50 \text{ kg } \underline{\text{n } \text{C}_{5}} \cdot \frac{1 \text{ kmol}}{72.5 \text{ kg}} = 0.6897 \text{ kmol } \text{n } \text{C}_{5} | \stackrel{5}{=} 0.445 \text{ kmol} = 0.5499 \text{ kmol} | z = 0.555 \text{ z}^{4}$$

DePriester Chart K_{C4} 2.05, K_{C5} 0.58, Result similar if use Raoult's law.

$$\frac{V}{V} = 0.555 = 0.445 = 1.3214 - 0.424 - 0.8976$$

F $0.58 + 1 - 0.05$
Check $f_{-} = \frac{-105 - 555}{V} = -\frac{42 - 445}{V} = 0.3000 - 29999 = 0.0K$
F $1 - 0.5 - 8976 = 1.42, 8976$
Eq. 3.23 $x_{C4} = -\frac{-105 - 555}{V} = 0.2857$, $1.42, 8976$
Eq. 3.23 $x_{C4} = -\frac{-105 - 555}{V} = 0.2857$, $1.42, 8976$
Eq. 3.23 $x_{C4} = -\frac{-105 - 555}{V} = 0.2857$, 0.443
b) From problem 2.D.g., K_{C4} 1.019 and K_{C5} 0.253.
Solving RR equation,
 $\frac{V}{F} = -\frac{-2.4}{V} = \frac{2F_{0-}}{V} = -\frac{-555}{0.2445} = 0.445$
 $x_{C5} = 7.143 y_{C4} - K_{C4} x_{C4} = 0.5857$, $y_{C5} = 0.4445$
 $x_{C5} = 7.143 y_{C4} - K_{C4} x_{C4} = 0.253 + 0.019$
NOT possible. Won't flash at 0°C.
2.D32. New problem in 4^{th} US edition. Was 2.D28 in 3^{rd} International Edition.
 $\frac{113}{V} = \frac{1}{2/3} + \frac{1}{2}$ Slopel 2 Through y x z_A 0.6 See figure
a.L
 $\frac{1}{F} = 33.33, x = 0.375$ (from Figure) $V = \frac{2}{2}F = 66.67$, $y = 0.72$ (from Figure)
 $\frac{V}{Fl_{B}} = 0.4 V_{A} = 0.466.67 - 26.67$, $L_{B} = 0.6F_{B} = 0.666.67 - 40.00$
b. $z_{C} = x_{A} = 0.375, -x_{C} = 0.15, -F_{C} = L_{A} = 33.33, From equilibrium y_{C} = 0.51$
 $\frac{V}{V} = \frac{1}{V} = 0.47 x_{A} = 0.466.7 - 26.67$, $L_{B} = 0.6F_{B} = 0.666.67 - 40.00$
b. $z_{C} = x_{A} = 0.375, -x_{C} = 0.15, -F_{C} = L_{A} = 33.33, From equilibrium y_{C} = 0.51$
 $\frac{V}{V} = \frac{1}{V} = 0.47 x_{A} = 0.67 x_{A} = 0.47 x_{A} = 0.47$

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/m³, liquid avg MW = 18.43, vapor density = 0.85 kg/m³, and

vapor avg MW = 26.02, T_{drum} = 94.1 $^{\circ}$ C, and Q = 6240.85 kW.

The diameter of the vertical drum in meters (with uperm in ft/s) is

 $D = \{ [4(MWv) V] / [3600 \pi \rho v u_{perm} (1 m/3.281 ft)] \}^{0.5} = \{ [4(26.02)(500)] / [3600(3.14159)(0.85)(1/3.281)u_{perm}] \}^{0.5}$ $F_{1v} = (WL/Wv) [\rho v / \rho L]^{0.5} = (9212.78/13010.57) [0.85/916.14]^{0.5} = 0.02157$

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} at 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/m³

 $D = \{ [4(MWv) V] / [3600 \pi \rho v u_{perm} (1 m/3.281 ft)] \}^{0.5} = \{ [4(26.14)(500)] / [3600(3.14159)(6.37)(1/3.281)u_{perm}] \}^{0.5}$

 $F_{lv} = (W_L/W_V) {\left[\rho v / \rho L\right]}^{0.5} = (9155.07/13068.27) {\left[6.37/836.89\right]}^{0.5} = 0.06112$

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/m³, MW_v = 26.10

 $D = \{ [4(MWv) V] / [3600 \pi \rho v u_{perm} (1 m/3.281 ft)] \}^{0.5} = \{ [4(26.10)(500)] / [3600(3.14159)(2.83)(1/3.281)u_{perm}] \}^{0.5} \\ 0.5 \\ 0.5 \\ F_{lv} = (Wt/Wv) [\rho v / \rho L] = (9173.91/13049.43) [2.83/874.58] = 0.0400$

Resulting Kvertical = .441162, uperm = 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/m³, $MW_V = 26.07$.

 $D = \{ [4(MWv) V] / [3600 \pi \rho v u_{perm} (1 m/3.281 ft)] \}^{0.5} =$ $\{ [4(26.07)(500)] / [3600(3.14159)(1.69)(1/3.281)u_{perm}] \}^{0.5}$ 0.5 D.5 = 0.5 $F_{lv} = (WL/Wv) [\rho v / \rho L] = (9188.82/13034.53) [1.69/893.99] = 0.0307$

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

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

In this case there is an advantage operating at a somewhat elevated pressure.

- This problem was 2.D13 in the 2nd edition of *SPE*. 2.E2.
 - 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, and y_{C6} y_{C8} K_{C6} x_{C6} K_{C8} x_{C8} 1.0 $X_{C6} K_{C6} K_{C8}$

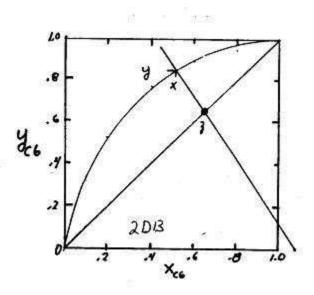
Substitute for xc6

Pick T, find Kc6 and Kc8 (e.g. from DePriester charts), solve for xc6. Then yc6 = Kc6xc6

T°C	KC6	Kc8	XC6	yc6 = Kc6 xc6
125	4	1.0	0	0
120	3.7		.0357	.321
110	3.0		.1379	.141
100	2.37		.2595	.615
90	1.8		.4406	.793
80	1.4		.650	.909
66.5	<u>1.0</u>	<u>.1</u> 7	1.0	1.0

Op Line Slope L 1 V/F .6 1.5, Intersection y = x = z = 0.65. V V/F .4

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.

 $\overline{\text{MW}}_{\text{L}}$ x_{C8} MW c₆ x_{C8} MW c₈ .52 86.17 .48 114.22 99.63 $\frac{114.22}{V^{L}} \frac{114.22}{145.98} \text{ ml/mol}$ c8 .659 .703 . C6 lbm L

 V_L 145.98 454 g/lbm ft

Now we can determine flow rates

$$V \frac{V}{F}F$$
 .4 10,000 4000 lbmol/h
W_v V MW v4000 90.38 361,520

L F V 6000 lbmol/h, W_L L MW_L 6000 99.63 597,780 lb/h

lb/h

$$u_{Perm} 3600 v$$
 6.3 3600 0.19135
D $\sqrt{4A_{Cs}/48333}$ /10.3 ft. Use 10.5 ft.

L ranges from $3 \times 10.5 = 31.5$ ft to $5 \times 10.5 = 52.5$ ft.

Note: This u_{Perm} is at 85% of flood. If we want to operate at lower % flood (say 75%) $u_{Perm 75\%} 0.75/0.85 u_{Perm 85\%} 0.75/0.85 .63 5.56$ Then at

75% of flood, $A_{Cs} = 94.44$ which is D = 10.96 or 11.0 ft.

2.E3. New problem 4^{th} 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, $L = \frac{L}{(MW)_{vapor}}$, but x and y are not known the molecular weights are unknown.

In weight units, V = F(V/F) = 2000 kg/h (0.35) = 700 kg/h. L = F – V = 1300 kg/h. In weight units the equilibrium data (Table 2-7) can be converted as follows:

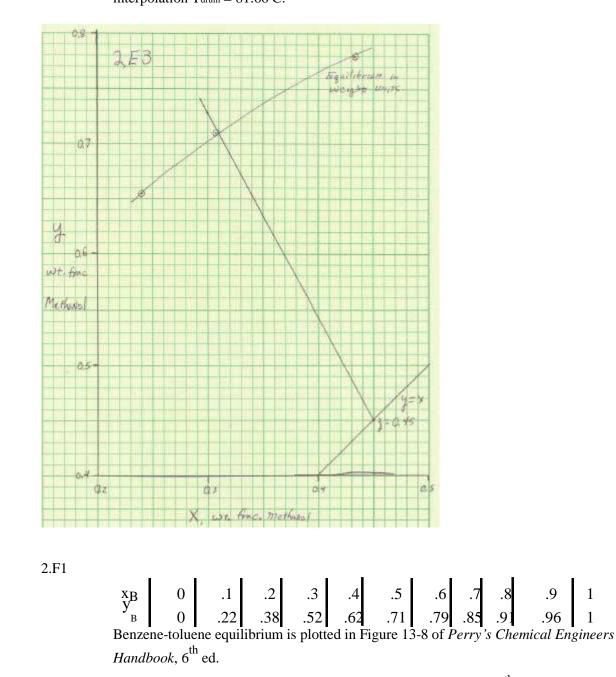
Basis: 1 mol, x = 0.4 and y = 0.729, T = 75.3 C

Liquid: 0.4 mol methanol \times 32.04 g/mol = 12.816 g

0.6 mol water × 18.016 g/mol = $\frac{10.806 \text{ g}}{\text{Total} = 23.622 \text{ g}} \rightarrow x = 0.5425 \text{ wt frac methanol}$

Vapor: 0.729 mol methanol = 23.357 g 0.271 mol water = $\underline{4.881 \text{ g}}$ 28.238 g \rightarrow y = 0.8271 wt frac methanol. Similar calculations for: 0.3 mole frac liquid give $x_{wt} = 0.433$ and $y_{wt} = 0.7793$, T = 78.0 C 0.2 mole frac liquid give $x_{wt} = 0.3078$ and $y_{wt} = 0.7099$, T = 81.7 C 0.15 mole frac liquid give $x_{wt} = 0.2389$ and $y_{wt} = 0.6557$, T = 84.4 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

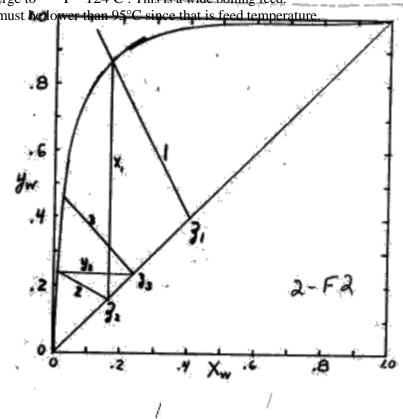
solve the problem, the scale could be increased resulting in better accuracy. By linear interpolation $T_{drum} = 81.66 \text{ C}$.



See Graph. Data is from *Perry's Chemical Engineers Handbook*, 6th ed., p. 13-12. 2.F2.

1

2.F3. Bubble Pt. At P = 250 kPa. Want
$$K_1 z_1$$
 1 .Solution uses DePriester chart for K values.
Guess T 18 C, K_1 1, K_2 .043, K_3 .00095, .52
Converge to T 0 C
 $\frac{Z_1}{K_1 1.0}$
Try T 0 C, K_1 1.93, 2 0.11, K_3 0.0033, 120.26



 $T_{d,1}$ 70 C : K_1 7.8, K_2 1.07, K .083 Guess V F 0.5 . Rachford Rice Eq.
 7.8
 1
 .517
 .07
 .091
 .083
 1
 .392

 f V F
 1
 6.8
 .5
 1
 .07
 .5
 1
 .083
 1
 .5
 .14
 V/F .6 gives f .6 .101 By linear interpolation: V/F .56. f 0.56 .0016 which is close enough for first trial. V V H/F 56. L 44 $x_i = \frac{z_i}{1 - K_i - 1 - V - F}$ and $y_i = K_i x_i$ x₁.1075 x₂ .088 x₃ .806 x 1.001 $y_1 .839 \hspace{0.1 in} y_2 \hspace{0.1 in} .094 \hspace{0.1 in} y_3 \hspace{0.1 in} .067 \hspace{0.1 in} y \hspace{0.1 in} .9999 \\ T25 \hspace{0.1 in} C \hspace{0.1 in} . \hspace{0.1 in} (Perry's \hspace{0.1 in} 6^{th} \hspace{0.1 in} ed; \hspace{0.1 in} p. \hspace{0.1 in} 3\text{-} 127), \hspace{0.1 in} and \hspace{0.1 in} (Perry's \hspace{0.1 in} 6^{th} \hspace{0.1 in} ed; \hspace{0.1 in} p. \hspace{0.1 in} 3\text{-} 138)$ Data: Pick 1 81.76 cal/g 44 3597.44 kcal/kmol 2 87.54 cal/g 72 6302.88 kcal/kmol 3 86.80 cal/g 114 9895.2 kcal/kmol T 0 C, C_{pL1} 0.576 cal / (g C) 44 25.34 kcal/(kmol C). at T 20 to 123 C, C_{pL3} 65.89 kcal/(kmol C) For T 75 C, C_{pL2} 39.66 kcal/(kmol C). (Himmelblau/Appendix E-7) at C_{pv} a bT cT² $\begin{array}{c} c_{pv} & a \ b1 \ c1 \\ propane & a = 16.26 \quad b = 5.398 \times 10^{-2} \\ n-pentane & a = 27.45 \quad b = 8.148 \times 10^{-2} \\ **n-octane & a = 8.163 \quad b = 140.217 \times 10^{-3} \end{array} \qquad \begin{array}{c} c = -3.134 \times 10^{-5} \\ c = -4.538 \times 10^{-5} \\ c = -44.127 \times 10^{-6} \end{array}$ ** Smith & Van Ness p. 106 Energy Balance: $E(T_d) = VH_v + Lh_L - Fh_F = 0$ Fh 100 .577 25.34 .091 39.66 .392 65.89 95.25 297,773 kcal/h .1075 25.34 .088 39.66 .806 65.89 70.25 117, 450 Lh 44 $V\!H_{v}_{_{0.94}\,_{6302.88}\,_{27.45}\,_{8.148}\,_{10^2}\,_{45}}^{}3597.4\,\,16.26\,\,5.398\,\,{10}^2$ 45 0.67 9895.3 8.163 140.217 10³ 45240,423 E T_{drum} 60,101 Thus, T_{drum} is too high. Converge on T 57.2 C: K 6.4, K 2.8, K 3.054

First Trial: Guess

VF 0.513, f 0.513 0.0027. V 51.3, L 48.7 For

x ₁ .137,	x ₂ .101,	x3	.762, x ₁	1.0000
y ₁ .878,	y ₂ .081,	y 3	.041, y ₁	1.0000

Fh_F 297,773; Lh_L 90,459; VH_v 209,999; E T_{drum} 2685

Thus T_{drum} must be very close to 57.3°C.

x₁.136, x₂.101, x₃.762, y₁.328, y₂.081, y₃.041 V 51.3

kmol/h, L 48.7 kmol/h

Note: With different data Tdrum 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 kJ/mol. For an adiabatic flash, h_F = [VHv + LhL]/F Vapor and liquid are in equilibrium. Saturated steam at 100 kPa is at T = 372.76K, h_L = 7.5214 kJ/mol, Hv = 48.19 kJ/mol
Mass balance: F = V + L where F in kmol/min = (1500 kg/min)(1 kmol/18.016 kg) = 83.259 kmol/min EB: Fh_F = VHv + Lh_L → (83.259 kmol/min)(17.604 kJ/mol)(1000 mol/kmol) = (48.19)(1000)V + (7.5214)(1000)L. Solve equations simultaneously. L = 62.617 kmol/min = 1128.12 kg/min and V = 20.642 kmol/min =

371.88 kg/min

2.G1. Used Peng-Robinson for hydrocarbons.

Find T_{drum} 33.13 C, L 34.82 and V 65.18 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 4^{th} edition. Part a. p = 31.26 kPa with V/F)_{feed} = 0.0009903. Part b. Use p_{feed} = 31.76 kPa, V/F)_{feed} = 0.0

Part c. Drum p = 3.9 bar, $T_{drmu} = 19.339$, V/F = 0.18605,

Liquid mole fractions: C1 = 0.14663, C2 = 0.027869 ($\sum = 0.05253$ is in spec), C5 = 0.6171, C6 = 0.3404. Vapor mole fractions: C1 = 0.68836, C2 = 0.20057, C5 = 0.9523, and C6 = 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:

	Х	у	K
Methane	0.004599	0.27039	58.79
n-butane	0.44567	0.52474	1.1774
n-pentane	0.54973	0.20488	0.37269

V/F)drum = 0.43419; V/F)feed = 0.3654; Q = -3183.4 cal/s

2.G4.

COMP	x(I)	y(I)
METHANE	0.12053E-01	0.84824
BUTANE	0.12978	0.78744E-01
PENTANE	0.29304	0.47918E-01
HEXANE	0.56513	0.25101E-01
V/F = 0.58354		

2.G5. N. Used NRTL. T = 368.07, Q = 14889 kW, 1^{st} liquid/total liquid = 0.4221,

Comp Liquid 1. x ₁ Liquid 2. x ₂ Vapor. y				
	Comp	$\mathbf{I}_{\mathbf{I}}$	$1 \cdot 1(111)(1 \cdot 2 \cdot \mathbf{X})$	Vapor V

Furfural	0.630	0.0226	0.0815
Water	0.346	0.965	0.820
Ethanol	0.0241	0.0125	0.0989

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.

COMP	x(I)	y(I)
METHANE	0.000273	0.04959
BUTANE	0.18015	0.47976
PENTANE	0.51681	0.39979
HEXANE	0.30276	0.07086
V/F = 0.40001		

2.H1. New Problem 4th ed A. 563.4 R, b.V/F = .4066. c. 18.264 psia

2.H3. New Problem. 4^{th} 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, 4^{th} ed. Answer: pdrum = 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 2H5. *New problem* 4th *ed*.

- 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. $T = 34.48^{\circ}C$
- d. $T = -1.586^{\circ}C$ and V/F = 0.0567

2H6. New problem in 4th ed.

4	C2.H6	

_										
1	А	В	С	D	E	F	G	Н	18	J
1		14								
2	Example	2-2 on spre	adsheet							
3	K1	7	K2	2.4	К3	0.8	K4	0.3		
4	z1	0.3	z2	0.1	z3	0.15	z4	0.45		
5	Guess V/	F 0.500823								
6	x1	0.074908	x2	0.058784	х3	0.166697	x4	0.692922		
7	y1	0.524353	y2	0.141081	уЗ	0.12	y4	0.207877	sum ↓	
8	yk-xi	0,449445		0.082297		-0.0467		-0.48505	-2.1E-07	
9								chk	-0.00021	
10		Goal seek	9I to zero	by changin	g B5				019	
4.4										