9th August 2010

P1. [20 marks, 2 marks for part (a), 4 marks for (b), 6 marks for (c) and 8 marks for (d)]

(a) Convert density in lb_m/ft^3A to kg/m³ = 85.3/(2.2*2.832*10⁻²)= 1369.1 kg/m³ Volume of pond = 50*15*2 = 1,500 m³. Therefore the mass of the pond is= **2.05356*10⁶ kg** Weight = 9.81*2.05356*10⁶ = **20.15*10⁶ N** (b)



If US gallons were used the answer would be 5770 planes.

(i) 200 crystal/min.mm; 10 crystals/min.mm².
(ii)
$$r = \frac{200 \text{ crystals} \quad 0.050 \text{ in} \quad 25.4 \text{ mm}}{\min \cdot \text{mm}} - \frac{10 \text{ crystals} \quad 0.050^2 \text{ in}^2 \quad (25.4)^2 \text{ mm}^2}{\min \cdot \text{mm}^2}$$

$$= 238 \text{ crystals} / \min \Rightarrow \frac{238 \text{ crystals} \quad 1 \text{ min}}{\min \quad 60 \text{ s}} = \frac{4.0 \text{ crystals} / \text{s}}{10 \text{ crystals} / \text{s}}$$

(iii)

$$D(\text{mm}) = \frac{D'(\text{in})}{|1|} = 25.4 \text{ mm} = 25.4 D'; \ r\left(\frac{\text{crystals}}{\text{min}}\right) = r' \frac{\text{crystals}}{|1|} \frac{60 \text{ s}}{|1|} = 60r'$$
$$\Rightarrow 60r' = 200(25.4D') - 10(25.4D')^2 \Rightarrow r' = 84.7D' - 108(D')^2$$

P2. [20 marks, 3 marks each for parts (a) and (b), 6 marks for (c) and 8 marks for (d)] (a)

(i) $2600*1.934*10^{-2}$ =**50.284 psi**. (ii) (275/33.9)*101.325=822kPa, (iii) $(3.00*1.01325*10^5)/(100^2) = 30.40 \text{ N/cm}^2$, (iv) 1atm - (20*10/760) = 0.737 atm, (v) 25.0*51.71= **1292.75 mm Hg (gauge)**, (vi) (25.0+14.696)*51.71= **2052.68 mm Hg (absolute).** (b) C T = -10 °F to °R, °C, K, T = -10°F+459.7=449.7°R/1.8=249K-273=-23°C (i) $\Delta T = 85 \ ^{\circ}\text{C} = 85 \text{K}, = 85 \ ^{\circ}\text{C} * 1.8 = 153 \ ^{\circ}\text{F} = 153 \ ^{\circ}\text{R},$ (ii) $\Delta T = 150 \ ^{\circ}\text{R} = 150 \ ^{\circ}\text{F}/1.8 = 83.3 \ ^{\circ}\text{C} = 83.3 \text{K}$ (iii) (c) (i) $(P' + 14.32) \times V' = 10.45n'(T' + 459.4)$ (ii) **0.316 lb-mole** of gas and **2.65 l_{bm}** of CO

(iii) **2735.5°F**. The ideal gas equation is an approximation of the real behavior of gases where several simplifying assumptions are made (no interactions between the molecules, no interaction with the surface, etc.). The ideal gas equation does not apply at high temperatures and pressures when the conditions approach the critical point.

							n n
1		C 1 1	1	D	· · ·	1 0 0 0 1 /	1
10	A daumntiona.	Stondord	aanditiana	1 Ion offst	of wotor 1	$\alpha + 1000 ka/m$	
	I ASSILLIDHOUS	Manualu	COLICITIONS		on water i	S I UNNI K9/III	
14	/ issumptions.	Standard	conditions,	Denoney	or mater i	$1,000 \text{ m}_{\text{m}}/\text{m}_{\text{m}}$	
· · ·	· 1		,	2		ý U	/

		Al	bu Dhabi		Kharg Island			
	MW	PPM (mg/L)	Weight F	Mol F	PPM (mg/L)	Weight F	Mol F	
Chloride (Cl)	35.453	24,300	0.55	0.50	23,000	0.56	0.50	
Sulfate (SO ₄)	96.064	3,420	0.08	0.03	3,100	0.07	0.02	
Bicarbonate (HCO ₃)	61	200	0.005	0.002	171	0.004	0.002	
Carbonate (CO ₃)	60	0	0	0	24	0.001	0.0003	
Sodium (Na)	23	13,900	0.32	0.44	13,044	0.32	0.44	
Magnesium (Mg)	54.94	1,560	0.04	0.02	1,500	0.04	0.02	
Calcium (Ca)	40.08	600	0.014	0.01	520	0.013	0.01	
SG		1.029			1.031			
No. of moles				1,372			1,292	
TDS		43,980			41,359			
Weight %		4.27%			4.01%			

P3. [25 marks, 8 marks each for parts (a) and (b), 6 marks for (c) and 3 marks for (d)]

 $\begin{array}{r} 2C_2H_4+O_2 \rightarrow 2C_2H_4O\\ C_2H_4+3O_2 \rightarrow 2CO_2+2H_2O\\ (a) \text{ Basis: 100 mol fed to reactor/h} \Rightarrow 25 \text{ mol O2/h, 75 mol C2H4/h} \end{array}$



(c)

$$\frac{\text{Scale factor}}{\text{h}} = \frac{2000 \text{ lbm C}_2\text{H}_4\text{O}}{\text{h}} \frac{|1 \text{ lb} - \text{mole C}_2\text{H}_4\text{O}|}{|44.05 \text{ lbm C}_2\text{H}_4\text{O}|} \frac{\text{h}}{|13.5 \text{ mol C}_2\text{H}_4\text{O}|} = 3.363 \frac{\text{lb} - \text{mol/h}}{\text{mol/h}}$$
$$n_{\text{C2H4}} = (3.363)(15.0) = \frac{50.4 \text{ lb} - \text{mol C}_2\text{H}_4\text{/h}}{|13.5 \text{ mol C}_2\text{H}_4\text{O}|} = \frac{3.363 \frac{\text{lb} - \text{mol/h}}{\text{mol/h}}}{|13.5 \text{ mol C}_2\text{H}_4\text{O}|}$$

P4. [15 marks, 6 marks parts (a) and 9 marks for part (b)]

- (a)
- (i) $P_{\text{final}} = 243 \text{ mm Hg}$. Since liquid is still present, the pressure and temperature must lie on the vapor-liquid equilibrium curve, where by definition the pressure is the vapor pressure of the
- (ii) Assuming ideal gas behavior for the vapor,

$$m(vapor) = \frac{(3.000 - 0.010) L}{(30 + 273.2) K} \frac{mol \cdot K}{0.08206 L \cdot atm} \frac{243 mm Hg}{760 mm Hg} \frac{1 atm}{mol} \frac{119.39 g}{mol} = 4.59 g$$

$$m(liquid) = \frac{10 mL}{1.489 g} \frac{1.489 g}{mL} = 14.89 g$$

$$m_{total} = m(vapor) + m(liquid) = \underline{19.5 g}$$

$$x_{vapor} = \frac{4.59}{19.48} = \underline{0.235 g vapor / g total}$$

- (b) Ethyl acetate has a vapour pressure of 118.3 mm Hg at 29.5°C and a normal boiling point of 77.0°C. Estimate the vapour pressure at 45°C using
 - (i) the Antoine equation:

$$\log_{10} p^* = 7.09808 - \frac{1238.71}{45 + 217} = 2.370 \Rightarrow p^* = 10^{2.370} = \frac{234.5 \text{ mm Hg}}{234.5 \text{ mm Hg}}$$

(ii)

$$\ln p^{*} = -\frac{\Delta \hat{H}_{v}}{R} \frac{1}{T} + B \Rightarrow -\frac{\Delta \hat{H}_{v}}{R} = \frac{\ln(p_{2}^{*}/p_{1}^{*})}{\frac{1}{T_{2}} - \frac{1}{T_{1}}} = \frac{\ln(760/118.3)}{(77.0+2732)K} = -4151K$$

$$B = \ln(p_{1}^{*}) + \frac{\Delta \hat{H}_{v}/R}{T_{1}} = \ln(118.3) + \frac{4151}{(29.5+273.2)K} = 18.49$$

$$\ln p^{*}(45^{\circ}C) = -\frac{4151}{(45+273.2)} + 18.49 \Rightarrow \underline{p^{*} = 231.0} \text{ mm Hg}$$

$$\frac{2310-234.5}{234.5} \times 100\% = -\frac{1.5\% \text{ error}}{29.5-77}$$
(iii)

$$p^{*} = \left(\frac{118.3-760}{29.5-77}\right)(45-29.5) + 118.3 = \frac{327.7}{100} \text{ mm Hg}$$

$$\frac{327.7-234.5}{234.5} \times 100\% = \frac{39.7\% \text{ error}}{29.7\% \text{ error}}$$

P5. [20 marks, 8 marks part (a) and 12 marks for part (b)]

$$\begin{array}{l} C_2H_4(g) + \frac{1}{2} O_2(g) \rightarrow C_2H_4O(g) \\ C_2H_4 + 3O_2 \rightarrow & 2CO_2 + 2H_2O \end{array}$$

Basis: 2 mol C₂H₄ fed to reactor



(a)

 $\begin{array}{l} \hline \hline 10\% \ \text{yield} \Rightarrow n_5 = \underbrace{0.500 \ \text{mol} \ \text{C}_2\text{H}_4 \ \text{consumed}}_{1 \ \text{mol} \ \text{C}_2\text{H}_4\text{O}} = \underbrace{0.350 \ \text{mol} \ \text{C}_2\text{H}_4\text{O}}_{1 \ \text{mol} \ \text{C}_2\text{H}_4} = \underbrace{0.350 \ \text{mol} \ \text{C}_2\text{H}_4\text{O}}_{2 \ \text{mol} \ \text{C}_2\text{H}_4} \\ \hline \hline \text{C balance on reactor:} \ (2)(2) = (2)(150) + (2)(0.350) + n_6 \Rightarrow n_6 = \underbrace{0.300 \ \text{mol} \ \text{CO}_2}_{2} \\ \hline \hline \text{Water formed:} \ n_7 = \underbrace{0.300 \ \text{mol} \ \text{CO}_2}_{1 \ \text{mol} \ \text{CO}_2} = \underbrace{0.300 \ \text{mol} \ \text{H}_2\text{O}}_{1 \ \text{mol} \ \text{CO}_2} = \underbrace{0.300 \ \text{mol} \ \text{H}_2\text{O}}_{2 \ \text{mol} \ \text{CO}_2} \\ \hline \hline \text{O balance on reactor:} \ (2)(1) = 2n_4 + 0.350 + (2)(0.300) + 0.300 \Rightarrow n_4 = \underbrace{0.375 \ \text{mol} \ \text{O}_2}_{2} \\ \hline \hline \text{Overall C balance:} \ 2n_1 = n_6 + 2n_5 = 0.300 + (2)(0.350) \Rightarrow n_1 = \underbrace{0.500 \ \text{mol} \ \text{C}_2\text{H}_4}_{2} \\ \hline \hline \text{Overall O balance:} \ 2n_2 = 2n_6 + n_7 + n_5 = (2)(0.300) + (0.300) + (0.350) \Rightarrow n_2 = \underbrace{0.625 \ \text{mol} \ \text{O}_2}_{2} \\ \hline \hline \frac{\text{Feed stream:}}{\text{Recycle stream:}} 80.0\% \ \text{C}_2\text{H}_4, \ 55.6\% \ \text{O}_2 \\ \hline \hline \frac{\text{Reactor inlet:}}{\text{Reactor inlet:}} \ 66.7\% \ \text{C}_2\text{H}_4, \ 33.3\% \ \text{O}_2 \\ \hline \hline \frac{\text{Reactor outlet:}}{\text{Reactor outlet:}} \ 53.1\% \ \text{C}_2\text{H}_4, \ 13.3\% \ \text{O}_2, \ 12.4\% \ \text{C}_2\text{H}_4\text{O}, \ 10.6\% \ \text{CO}_2, \ 10.6\% \ \text{H}_2\text{O} \\ \hline \end{array}$

Mass of ethylene oxide =
$$\frac{0.350 \text{ mol } \text{C}_2\text{H}_4\text{O}}{1 \text{ mol}} \frac{44.05 \text{ g}}{10^3 \text{ g}} = 0.0154 \text{ kg}$$

(b) References for enthalpy calculations :C (s) , H_2 (g) , O_2 (g) at 25 °C

$$\begin{split} \hat{H}_i(T) &= \Delta \hat{H}_{fi}^\circ + \int_{25}^{T} C_p dT \text{ for } C_2 H_4 \\ &= \Delta \hat{H}_f^\circ + \int_{298}^{T+273} C_p dT \text{ for } C_2 H_4 O \\ &= \Delta \hat{H}_{fi}^\circ + \hat{H}_i \text{ (table B.8)} \\ &= \Delta \hat{H}_f^\circ \text{ for } H_2 O(1) \text{ for } O_2, \text{ } CO_2, \text{ } H_2 O(g) \end{split}$$

	verall Proce	ss		Reactor					
Substance	$n_{\rm in}$	$\hat{H}_{ ext{in}}$	n _{cut}	\hat{H}_{out}	substance	$n_{\rm in}$	\hat{H}_{in}	n _{out}	\hat{H}_{out}
Sustaire	(mol)	(kJ/mol)	(mol)	(kJ/mol)	suosuice	(mol)	(kJ/mol)	(mol)	(kJ/mol)
C ₂ H ₄	0.500	52.28	-	-	C ₂ H ₄	2	79.26	1.50	79.26
0 ₂	0.625	0	-	-	O2	1	13.37	0.375	13.37
C ₂ II ₄ O	-	-	0.350	-51.00	C_2II_4O	-	-	0.350	-19.99
CO ₂	-	-	0.300	-393.5	CO ₂	-	-	0.300	-374.66
$H_2O(l)$			0.300	285.84	$H_2O(g)$			0.300	226.72

<u>Energy balance on process</u>: $Q = \Delta H = \sum_{\text{out}} n_i \hat{H}_i - \sum_{\text{in}} n_i \hat{H}_i = \underline{-248 \text{ kJ}}$

Energy balance on reactor:
$$Q = \Delta H = \sum_{\text{out}} n_i \hat{H}_i - \sum_{\text{in}} n_i \hat{H}_i = \underline{-236 \text{ kJ}}$$

Bonus Problem [10 marks]

<u>void volume of bed:</u> 100 cm³ – (233.5 – 184)cm³ = 50.5 cm³ <u>porosity:</u> 50.5 cm³ void/184 cm³ total = 0.274 cm³ void/cm³ total <u>bulk density:</u> 600 g/184 cm³ = 3.26 g/cm³ <u>absolute density:</u> 600 g/(184 – 50.5)cm³ = 4.49 g/cm³