

Lecture 23

Chemical Reaction Engineering (CRE) is the field that studies the rates and mechanisms of chemical reactions and the design of the reactors in which they take place.

Web Lecture 23

Class Lecture 19-Thursday 3/21/2013

Multiple Reactions with **Heat Effects**



Multiple Reactions with Heat Effects

PFR/PBR:

$$\frac{dT}{dV} = \frac{Ua(T_a - T) + \sum_{i=1}^n (-r_{ij})(\Delta H_{Rxi j})}{\sum_{j=1}^m F_j C_{Pj}}$$

CSTR:


$$UA(T_a - T) - F_{A0} \sum_{j=1}^m C_{Pi} \theta_i (T - T_0) \sum_{i=1}^q (r_{ij}) H_{Rxi j}(T) = 0$$

These equations are coupled with the **mole balances** and **rate law** equations.

Multiple Reactions with Heat Effects

Multiple Reactions

Make sure each rate of reaction and rate law are in respect to the same species (example, A); Subscripts must agree

$$\frac{dT}{dV} = \frac{Q_g - Q_r}{\sum F_i C_{Pi}} \quad Q_g = \sum r_{ij} \Delta H_{Rxij} = r_{1A} \Delta H_{Rx1A} + r_{2A} \Delta H_{Rx2A}$$


Multiple Reactions with Heat Effects

Multiple Reactions

1) Mole Balances: – every species (no conversion!)

2) Rate Laws:

– relative rates

– net rates

3) Stoichiometry: $C_A = C_{T0} \frac{F_A}{F_T} p \frac{T_0}{T}$

$$\frac{dp}{dW} = \frac{-\alpha F_T T}{2p F_{T0} T_0}$$

Multiple Reactions with Heat Effects

Multiple Reactions

4) Heat Effects:

$$\frac{dT}{dV} = \frac{Q_g - Q_r}{\sum F_i C_{Pi}}$$

Q_g = heat produced

Q_r = heat removed

$$Q_g = \sum r_{ij} \Delta H_{Rxij} \quad (\text{must have matching } i, j)$$

$$Q_r = Ua(T - T_a)$$

Multiple Reactions with Heat Effects

4) Heat Effects:

$$\frac{dT}{dV} = \frac{Q_g - Q_r}{\sum F_i C_{Pi}}$$

$$Q_g = r_{1A} \Delta H_{R1A} + r_{2A} \Delta H_{R2A}$$

$$Q_r = Ua(T - T_a)$$

$$\sum F_i C_{Pi} = F_A C_{PA} + F_B C_{PB} + F_C C_{PC} + F_D C_{PD}$$

$$\frac{dT_a}{dV} = \frac{Ua(T - T_a)}{\dot{m}_i C_{Pcool}}$$

5) Parameters

$$E_1, E_2, F_{A0}, Ua, \dots \text{etc}$$

Multiple Reactions with Heat Effects

4) Heat Effects:

$$\frac{dT}{dV} = \frac{Q_g - Q_r}{\sum F_i C_{Pi}} \quad (23)$$

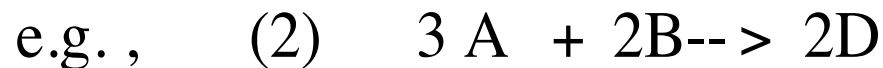
$$\Delta H_{Rx1A} = -12kJ / (\text{mole of A reacted in reaction 1})$$

$$\Delta H_{Rx2B} = +8kJ / (\text{mole of B reacted in reaction 2})$$

$$Q_g = r_{1A} \Delta H_{Rx1A} + r_{2B} \Delta H_{Rx2B}$$

Use relative rates of reaction

to get r_{2B} in terms of the rate law that is given for reaction 2,



$$-r_{2A} = k_{2A} C_A^3 C_B^2 \quad \text{then} \quad r_{2B} = \frac{2}{3} r_{2A}$$

The complex gas phase reactions



take place in a 10 dm³ PFR with a heat exchanger. Plot the temperature, concentrations, molar flow rates down the length of the reactor for the following operations. E.g., Note any maximums or minimums on your plot along with how they change for the different types of operations.

- Adiabatic operation
- Heat exchange with constant T_a
- Co current heat exchange
- Counter current heat exchange
- For parts (c) and (d), plot Q_r and Q_g down the length of the reactor. What do you observe?

Additional Information

$$C_{P_A} = 10 \text{ cal/mol/K}$$

$$C_{P_C} = 30 \text{ cal/mol/K}$$

$$C_{P_B} = 10 \text{ cal/mol/K}$$

$$C_{P_D} = 20 \text{ cal/mol/K}$$

$$k_{1A} = 40 \left(\text{dm}^3/\text{mol} \right)^2 / \text{s/K} \text{ at } 300\text{K} \text{ and } E_1 = 8,000 \text{ cal/mol}$$

$$k_{2C} = 2 \left(\text{dm}^3/\text{mol} \right)^2 / \text{s/K} \text{ at } 300\text{K} \text{ and } E_2 = 12,000 \text{ cal/mol}$$

$$C_{T_0} = 0.2 \text{ mol/dm}^3, \quad C_{P_{\text{Cool}}} = 1 \text{ cal/g/K}, \quad \dot{m}_{\text{Cool}} = 20 \text{ g/s}$$

$$Ua = 80 \text{ cal/dm}^3/\text{s/K}, \quad T_{a0} = 325\text{K}, \quad T_0 = 300\text{K}$$

$$F_{A0} = 5 \text{ mol/s}, \quad F_{B0} = 10 \text{ mol/s}, \quad F_{C0} = 0, \quad F_{D0} = 0$$

Calculated values of DEQ variables

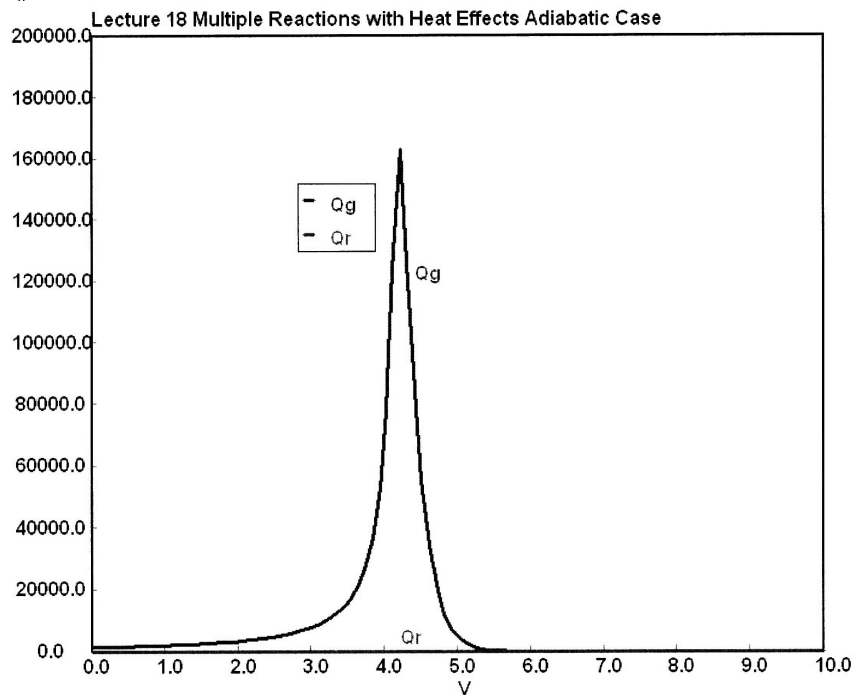
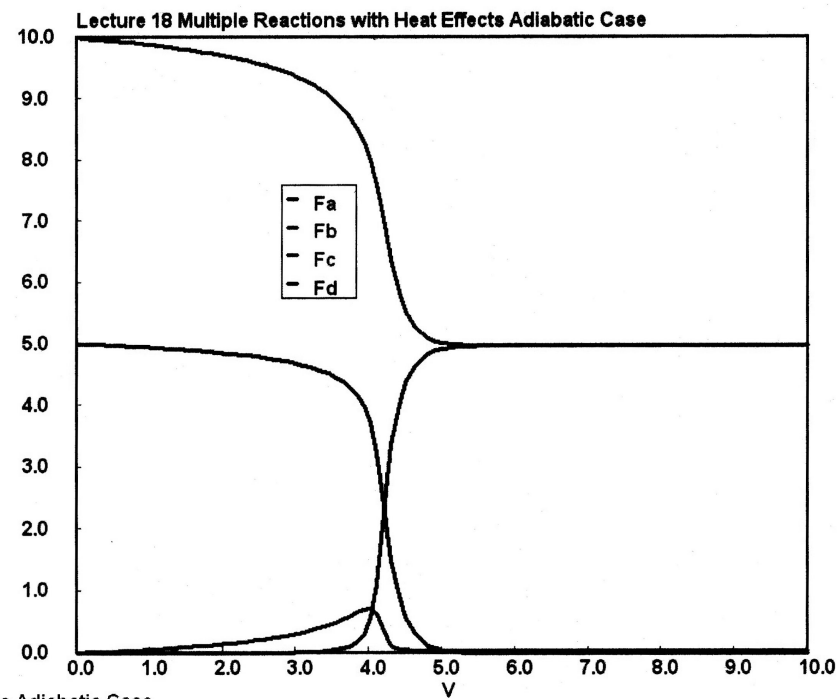
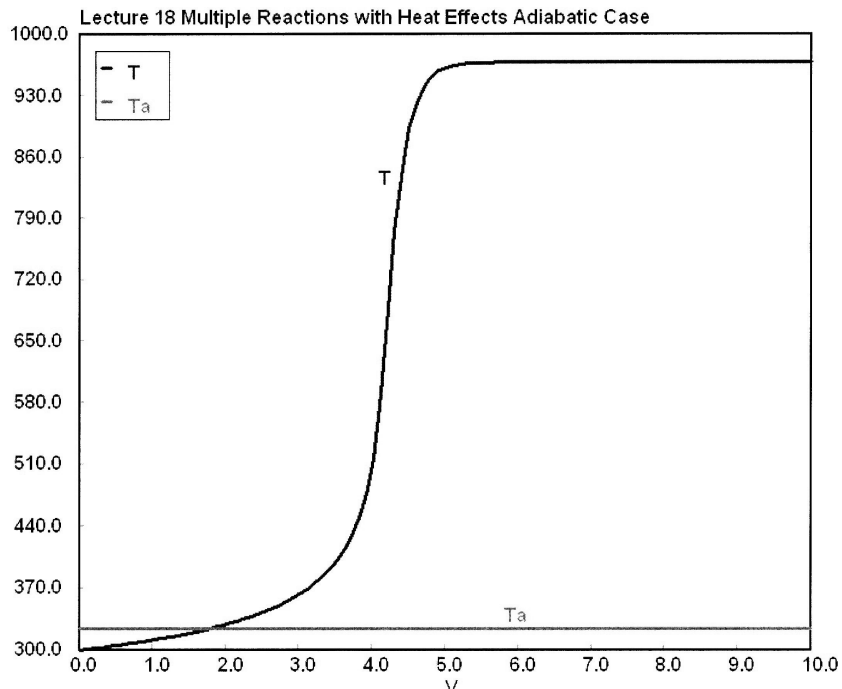
	Variable	Initial value	Final value
1	Ca	0.0666667	2.49E-05
2	Cb	0.1333333	0.0611818
3	Cc	0	0.0012689
4	Cpa	10.	10.
5	Cpb	10.	10.
6	Cpc	30.	30.
7	Cpco	10.	10.
8	Cpd	20.	20.
9	Cto	0.2	0.2
10	DH1b	-1.5E+04	-1.5E+04
11	DH2a	-10000.	-10000.
12	E1	8000.	8000.
13	E2	1.2E+04	1.2E+04
14	Fa	5.	0.0019942
15	Fb	10.	4.900364
16	Fc	0	0.1016299
17	Fd	0	4.896376
18	Ft	15.	9.900364
19	k1a	40.	6734.733
20	k2c	2.	4369.388
21	m	50.	50.
22	Qg	1422.222	20.21055
23	Qr	-2000.	1126.152
24	R	1.987	1.987
25	r1a	-0.0474074	-0.0006277
26	r1b	-0.0948148	-0.0012553
27	r1c	0.0474074	0.0006277
28	r2a	0	-0.000138
29	r2c	0	-0.000138
30	r2d	0	0.0002761
31	ra	-0.0474074	-0.0007657
32	rb	-0.0948148	-0.0012553
33	rc	0.0474074	0.0004896
34	rd	0	0.0002761
35	sumFiCpi	150.	150.
36	T	300.	485.4075
37	Ta	325.	471.3306
38	Ta55	325.	325.
39	To	300.	300.
40	Ua	80.	80.
41	V	0	10.
42	y	1.	1.

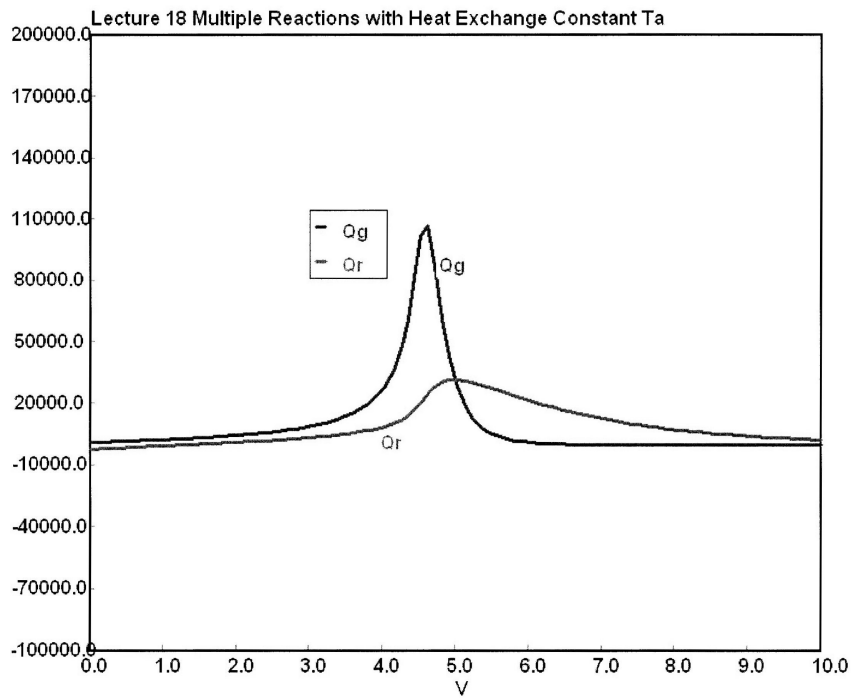
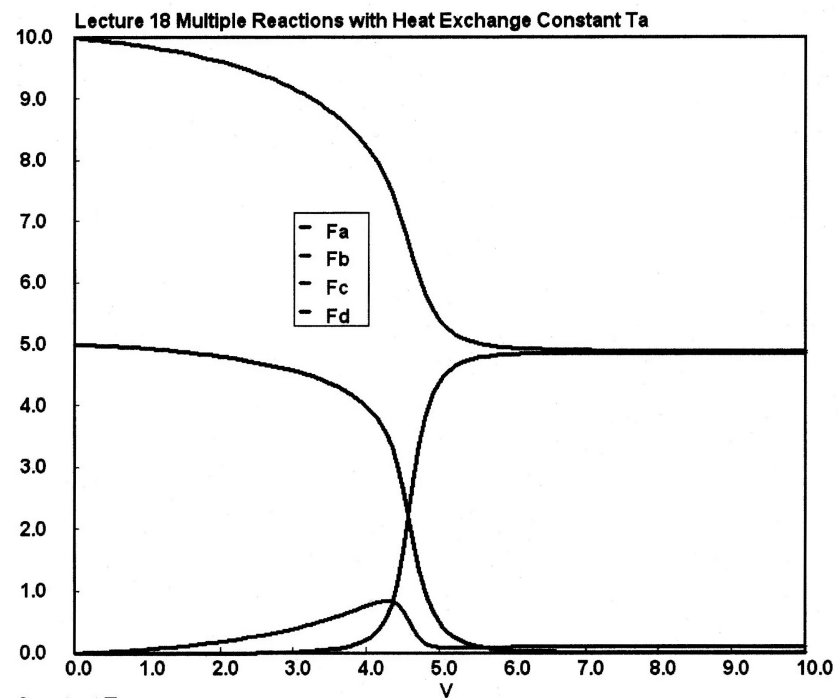
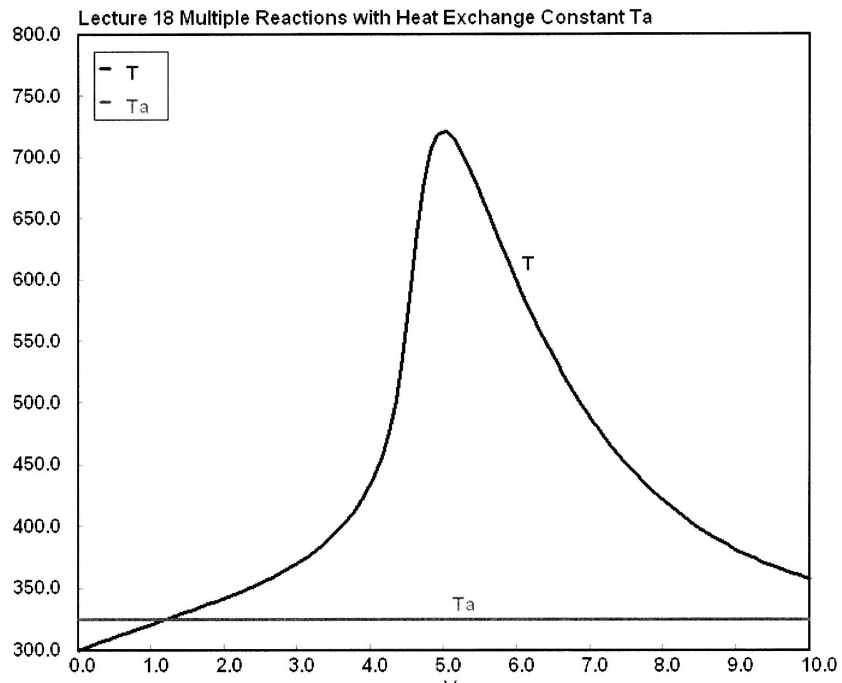
Differential equations

- $d(\text{Fa})/d(V) = \text{ra}$
- $d(\text{Fb})/d(V) = \text{rb}$
- $d(\text{Fc})/d(V) = \text{rc}$
- $d(\text{Fd})/d(V) = \text{rd}$
- $d(T)/d(V) = (\text{Qg}-\text{Qr})/\text{sumFiCpi}$
- $d(\text{Ta})/d(V) = \text{Ua}*(\text{T}-\text{Ta})/\text{m}/\text{Cpco}$

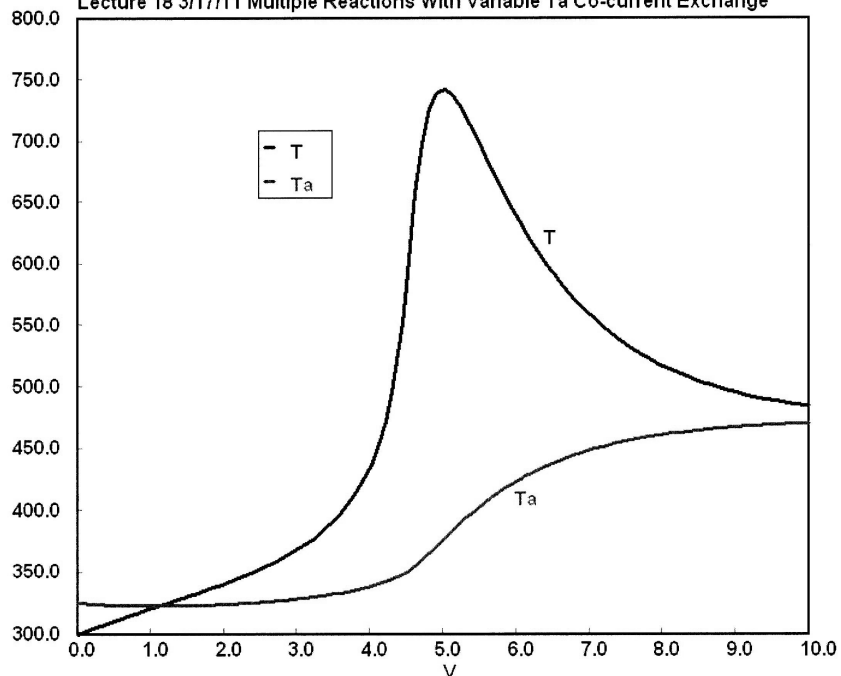
Explicit equations

- $E2 = 12000$
- $y = 1$
- $R = 1.987$
- $Ft = \text{Fa}+\text{Fb}+\text{Fc}+\text{Fd}$
- $To = 300$
- $k2c = 2*\exp((E2/R)*(1/300-1/T))$
- $E1 = 8000$
- $Cto = 0.2$
- $Ca = Cto*(\text{Fa}/\text{Ft})*(\text{To}/\text{T})^y$
-
- $Cc = Cto*(\text{Fc}/\text{Ft})*(\text{To}/\text{T})^y$
- $r2c = -k2c*Ca*Cc$
- $Cpco = 10$
- $m = 50$
- $Cb = Cto*(\text{Fb}/\text{Ft})*(\text{To}/\text{T})^y$
- $k1a = 40*\exp((E1/R)*(1/300-1/T))$
- $r1a = -k1a*Ca*Cb^2$
- $r1b = 2*r1a$
- $rb = r1b$
- $r2a = r2c$
- $\text{DH1b} = -15000$
- $\text{DH2a} = -10000$
- $r1c = -r1a$
- $\text{Ta55} = 325$
- $\text{Cpd} = 20$
- $\text{Cpa} = 10$
- $\text{Cpb} = 10$
- $\text{Cpc} = 30$
- $\text{sumFiCpi} = \text{Cpa}*\text{Fa}+\text{Cpb}*\text{Fb}+\text{Cpc}*\text{Fc}+\text{Cpd}*\text{Fd}$
- $\text{rc} = r1c+r2c$
- $\text{Ua} = 80$
- $r2d = -2*r2c$
- $\text{ra} = r1a+r2a$
- $\text{rd} = r2d$
- $\text{Qg} = r1b*\text{DH1b}+r2a*\text{DH2a}$
- $\text{Qr} = \text{Ua}*(\text{T}-\text{Ta})$

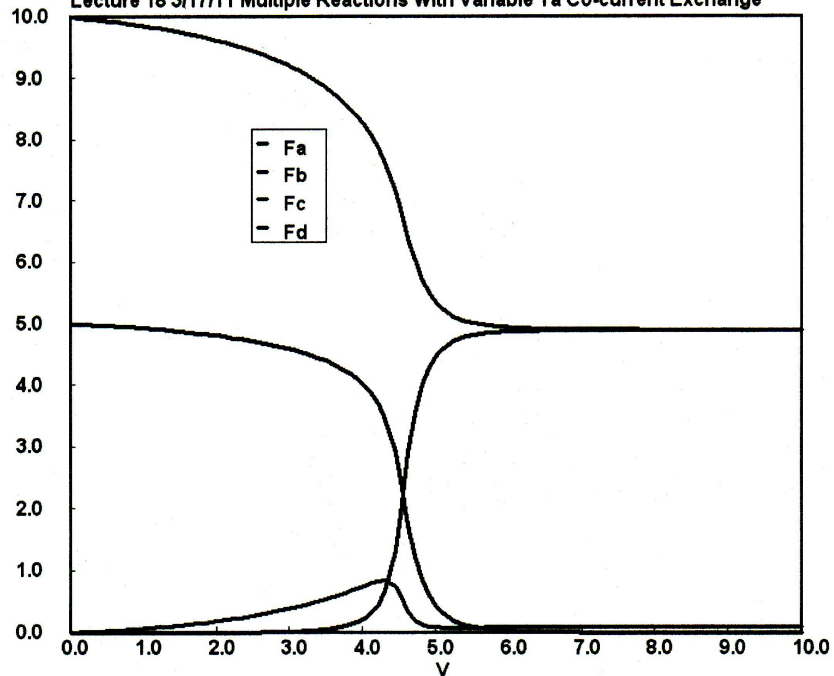




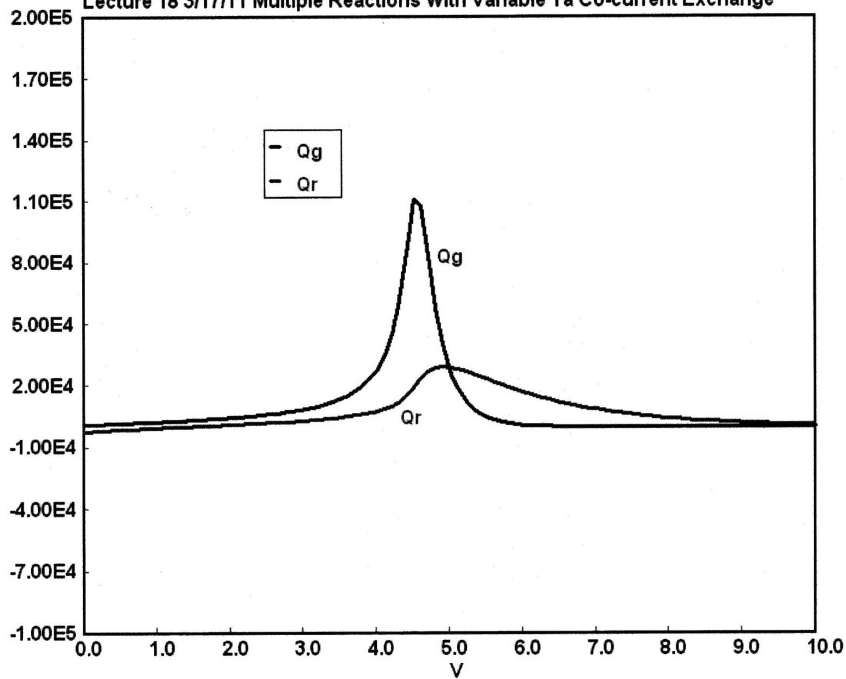
Lecture 18 3/17/11 Multiple Reactions With Variable Ta Co-current Exchange



Lecture 18 3/17/11 Multiple Reactions With Variable Ta Co-current Exchange



Lecture 18 3/17/11 Multiple Reactions With Variable Ta Co-current Exchange



Multiple Reactions with Heat Effects

Multiple Reactions

4) Heat Effects:
$$\frac{dT}{dV} = (-r_A)(-\Delta H_{Rx}) - \frac{Ua(T - T_a)}{\sum F_i C_{Pi}}$$

$$\frac{dT}{dV} = \frac{Q_g - Q_r}{\sum F_i C_{Pi}}$$

Q_g = heat produced

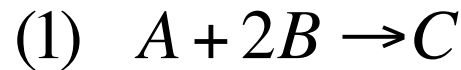
Q_r = heat removed

$$Q_g = \sum r_{ij} \Delta H_{Rij}$$

(must have matching i, j)

Multiple Reactions with Heat Effects in a PFR and CSTR

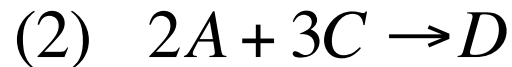
Examples:



$$-r_{1A} = k_{1A} C_A C_B^2$$

and

$$\Delta H_{R1A} = -20,000 \text{ cal/mol A}$$



$$-r_{2C} = k_{2C} C_A^2 C_C^3$$

and

$$\Delta H_{R2A} = 10,000 \text{ cal/mol A}$$

Example A: Liquid Phase CSTR



NOTE: The specific reaction rate k_{1A} is defined with respect to species A.



NOTE: The specific reaction rate k_{2C} is defined with respect to species C.

Example A: Liquid Phase CSTR

The complex liquid phase reactions take place in a 2,500 dm³ CSTR. The feed is equal molar in A and B with $F_{A0}=200$ mol/min, the volumetric flow rate is 100 dm³/min and the reaction volume is 50 dm³.

Find the concentrations of A, B, C and D existing in the reactor along with the existing selectivity.

Plot F_A , F_B , F_C , F_D and $S_{C/D}$ as a function of V

Example A: Liquid Phase CSTR

Solution

Liquid Phase CSTR

1) Mole Balances:

$$(1) \quad f(C_A) = v_0 C_{A0} - v_0 C_A + r_A V$$

$$(2) \quad f(C_B) = v_0 C_{B0} - v_0 C_B + r_B V$$

$$(3) \quad f(C_C) = -v_0 C_C + r_C V$$

$$(4) \quad f(C_D) = -v_0 C_D + r_D V$$

2) Net Rates:

$$(5) \quad r_A = r_{1A} + r_{2A}$$

Example A: Liquid Phase CSTR

3) Stoichiometry:

$$(16) \quad C_A = F_A / v_0$$

$$(17) \quad C_B = F_B / v_0$$

$$(18) \quad C_C = F_C / v_0$$

$$(19) \quad C_D = F_D / v_0$$

4) Parameters:

$$(20) \quad v_0 = 100 \text{ dm}^3/\text{min}$$

$$(21) \quad k_{1A} = 10 (\text{dm}^3/\text{mol})^2 / \text{min}$$

$$(22) \quad k_{2C} = 15 (\text{dm}^3/\text{mol})^4 / \text{min}$$

Example B: Liquid Phase PFR

Takes place in a PFR. The feed is equal molar in A and B and $F_{A0}=200$ mol/min and the volumetric flow rate is 100 dm³/min. The reaction volume is 50 dm³ and the rate constants are:

$$k_{1A} = 10 \left(\text{dm}^3 / \text{mol} \right)^2 / \text{min}$$

$$k_{2C} = 15 \left(\text{dm}^3 / \text{mol} \right)^4 / \text{min}$$

Rate laws are the same as Example A.

Plot F_A , F_B , F_C , F_D and $S_{C/D}$ as a function of V .

Example B: Liquid Phase PFR

1) Mole Balances:

$$(1) \quad \frac{dF_A}{dV} = r_A \quad (F_{A0} = 200 \text{ mol/min})$$

$$(2) \quad \frac{dF_B}{dV} = r_B \quad (F_{B0} = 200 \text{ mol/min})$$

$$(3) \quad \frac{dF_C}{dV} = r_C \quad V_F = 50 \text{ dm}^3$$

$$(4) \quad \frac{dF_D}{dV} = r_D$$

Example B: Liquid Phase PFR

2) Net Rates:

$$(5) \quad r_A = r_{1A} + r_{2A}$$

$$(6) \quad r_B = r_{1B}$$

$$(7) \quad r_C = r_{1C} + r_{2C}$$

$$(8) \quad r_D = r_{2D}$$

2) Rate Laws:

$$(9) \quad r_{1A} = -k_{1A} C_A C_B^2$$

$$(10) \quad r_{2C} = -k_{2C} C_A^2 C_C^3$$

Example B: Liquid Phase PFR

2) Relative Rates:

$$\frac{r_{1A}}{-1} = \frac{r_{1B}}{-2} = \frac{r_{1C}}{1} \quad \text{Reaction 1}$$

$$(11) \quad r_{1B} = 2r_{1A}$$

$$(12) \quad r_{1C} = -r_{1A} \quad \text{Reaction 2}$$

$$\frac{r_{2A}}{-2} = \frac{r_{2C}}{-3} = \frac{r_{2D}}{1}$$

$$(13) \quad r_{2A} = 2/3 r_{2C}$$

$$(14) \quad r_{2D} = -1/3 r_{2C}$$

Example B: Liquid Phase PFR

2) Rate Laws:

$$r_{1A} = -k_{1A} C_A C_B^2 \quad (5) \quad k_{1A} = k_{1A1} \exp\left[\frac{E_1}{R}\left(\frac{1}{T_1} - \frac{1}{T}\right)\right] \quad (6)$$

$$r_{2C} = -k_{2C} C_A^2 C_C^3 \quad (7) \quad k_{2C} = k_{2C2} \exp\left[\frac{E_2}{R}\left(\frac{1}{T_2} - \frac{1}{T}\right)\right] \quad (8)$$

$$r_A = r_{1A} + r_{2B} \quad (9) \quad r_B = r_{1B} \quad (10)$$

$$r_C = r_{1C} + r_{2C} \quad (11) \quad r_D = r_{2D} \quad (12)$$

$$r_{1C} = -r_{1A} \quad (13) \quad r_{1B} = 2r_{1A} \quad (13)$$

$$r_{2A} = 2/3 r_{2C} \quad (15) \quad r_{2D} = -1/3 r_{2C} \quad (16)$$

Example B: Liquid Phase PFR

3) Stoichiometry:

$$C_A = C_{T0} \frac{F_A}{F_T} p \frac{T_0}{T} \quad (17)$$

$$C_B = C_{T0} \frac{F_B}{F_T} p \frac{T_0}{T} \quad (18)$$

$$C_C = C_{T0} \frac{F_C}{F_T} p \frac{T_0}{T} \quad (19)$$

$$C_D = C_{T0} \frac{F_D}{F_T} p \frac{T_0}{T} \quad (20)$$

$$F_T = F_A + F_B + F_C + F_D \quad (21)$$

$$\frac{dp}{dV} = \frac{-\alpha\rho}{2p} \frac{F_T}{F_{T0}} \frac{T}{T_0} \quad (22)$$

Multiple Reactions with Heat Effects

4) Heat Effects:

$$\frac{dT}{dV} = \frac{Q_g - Q_r}{\sum F_i C_{Pi}} \quad (23)$$

$$\frac{dT_a}{dV} = \frac{Ua(T - T_a)}{\dot{m}_i C_{Pcool}} \quad (26) \quad \text{Co-Current}$$

$$Q_g = r_{1A} \Delta H_{R1A} + r_{2A} \Delta H_{R2A} \quad (24)$$

$$Q_r = Ua(T - T_a) \quad (25)$$

$$\sum C_P = F_A C_{PA} + F_B C_{PB} + F_C C_{PC} + F_D C_{PD} \quad (27)$$

Parameters:

$$E_1, E_2, F_{A0}, \dots$$

Selectivity

If one were to write $S_{C/D} = F_C / F_D$ in the Polymath program, Polymath would not execute because at $V=0$, $F_C=0$ resulting in an undefined volume (infinity) at $V=0$. To get around this problem we start the calculation 10^{-4} dm³ from the reactor entrance where F_D will not be zero and use the following IF statement.

$$(15) \quad S_{C/D} = \text{if } (V > 0.001) \text{ then } \left(\frac{F_C}{F_D} \right) \text{ else } (0)$$

Selectivity

3) Stoichiometry:

$$(16) \quad C_A = F_A / v_0$$

$$(17) \quad C_B = F_B / v_0$$

$$(18) \quad C_C = F_C / v_0$$

$$(19) \quad C_D = F_D / v_0$$

Parameters:

$$(20) \quad v_0 = 100 \text{ dm}^3 / \text{min}$$

$$(21) \quad k_{1A} = 10 (\text{dm}^3 / \text{mol})^2 / \text{min}$$

$$(22) \quad k_{2C} = 15 (\text{dm}^3 / \text{mol})^4 / \text{min}$$

End of Web Lecture 23

Class Lecture 18