

10.1 For the 4 schemes:  $G, N, N, G$   $\leftarrow$

10.3 Here we have:  $N, N, N, G$   $\leftarrow$

10.5 a) We should operate at the maximum allowable temperature when

$$E_1 > E_2 \text{ and } E_1 > E_3 \quad \leftarrow \quad \text{a)}$$

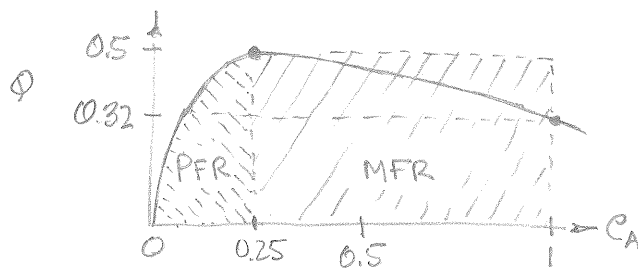
b) We should have a falling temperature when

$$E_1 > E_3 \text{ and } E_1 < E_2 \quad \leftarrow \quad \text{b)}$$

10.7 From Example 1

$$\phi(S/A) = \frac{0.2C_A}{0.025 + 0.2C_A + 0.4C_A^2}, \text{ and } \begin{cases} C_A = 0.25 \\ C_S = 0.375 \\ \phi_{\max} = \frac{0.375}{1 - 0.25} = 0.5 \end{cases}$$

The  $\phi$  vs  $C_A$  curve is



so use MFR followed by PFR



For MFR:  $C_{sm} = \phi(C_{A0} - C_A) = 0.5(1 - 0.25) = 0.375$

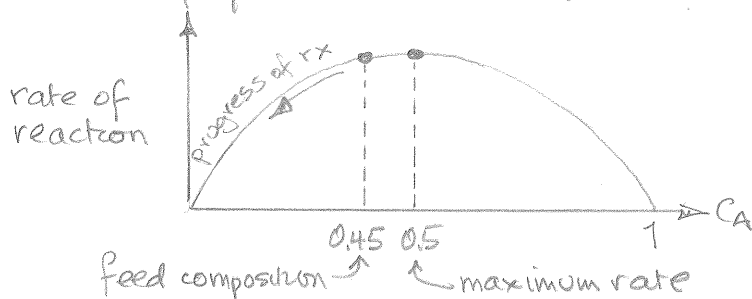
For PFR:  $C_{sp} = \int \phi dC_A = \int \frac{8C_A dC_A}{1 + 8C_A + 16C_A^2} = 8 \int_0^{0.25} \frac{C_A}{(1 + 4C_A)^2} dC_A$

But from a table of integrals:  $\int \frac{x}{(a+bx)^2} dx = \frac{a}{b^2(a+bx)} + \frac{1}{b^2} \ln(a+bx) \Big|_1^2$

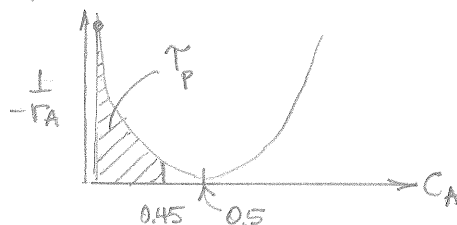
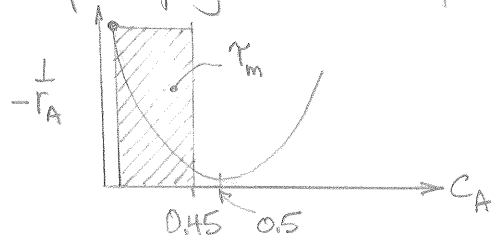
So  $C_{sp} = \left[ \frac{1}{16} \cdot \frac{1}{1+4x} + \frac{1}{16} \ln(1+4x) \right]_0^{0.25} = \frac{1}{2} \left[ \frac{1}{2} + \ln 2 - 1 \right] = 0.0966$

$\therefore C_{stabl} = C_{sm} + C_{sp} = 0.375 + 0.0966 = 0.4716 \quad \leftarrow$

10.9 Since  $C_A + C_B = 1$  for all  $C_A$  the maximum rate is at  $C_A = C_B = 0.5$ . Check this if you don't believe it. Thus



compare plug with mixed flow performance



Plug flow is best. Do not use recycle as recommended in Fig P9

10.11



The feed has  $C_R/C_T = \infty$ .

(a) So to maximize  $C_R/C_T$  do not react at all } a)  
 The design of Fig P11 is no good }

(b) To minimize  $C_R/C_T$  run to completion. All the R will disappear. b)

Next reaction 1 is second order } keep  $C_A$  low  
 reaction 2 is first order }

So use a large MFR, you will end up with most  $C_T$  b)

10.13

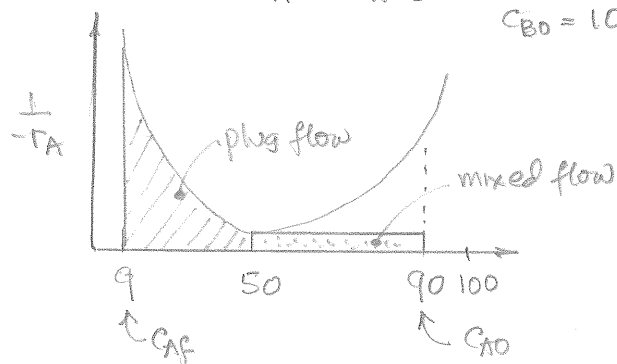


$$C_{A0} = 90$$

$$C_{B0} = 10$$

$$C_{Af} = 9$$

$$C_{Bf} = 91$$



Use a MFR followed by a PFR

10.15

Since  $E_1 > E_2$  use a high temperature, or  $T = 90^\circ\text{C}$

At  $90^\circ\text{C} = 363\text{K}$   $k_1 = 30 e^{-20000/8.314(363)} = 0.0397 \text{ min}^{-1}$

$k_2 = 1.9 e^{-15000/8.314(363)} = 0.0132 \text{ min}^{-1}$

$\therefore k_2/k_1 = 0.3322$

From Fig 8.13 or Eqs 8.8 & 8.9

$\frac{C_{R\max}}{C_{A0}} = 0.58$

$t_{\text{opt}} = \frac{\ln(k_2/k_1)}{k_2 - k_1} = \frac{\ln 0.3322}{0.0132 - 0.0397} = 41.6 \text{ min}$

10.17

Let  $A = \text{C}_3\text{H}_6$

$B = \text{O}_2$

$R = \text{C}_3\text{H}_4\text{O}$  (desired)

then



$k_2/k_1 = 0.1$

$k_3/k_1 = 0.25$

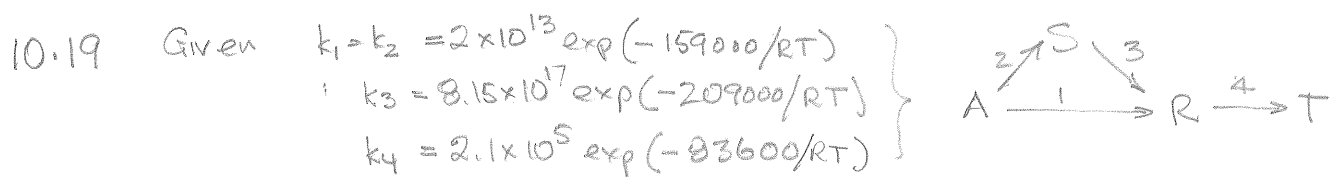
Rewrite the rates



From the Denbigh reaction set, eq 49 gives

$\frac{C_{R\max}}{C_{A0}} = \frac{k_1}{k_{12}} \left( \frac{k_{12}}{k_3} \right)^{k_3/(k_3-k_{12})} = \frac{10}{11} \left( \frac{11}{2.5} \right)^{2.5/(2.5-11)} = 0.588$

Use plug flow



Since  $E_4$  is much smaller than all the other  $E$  values use the highest allowable temperature, or 1200 K.

At 1200 K

$$k_1 = 2.4 \times 10^6 \text{ hr}^{-1}$$

$$k_2 = 2.4 \times 10^6 \text{ hr}^{-1}$$

$$k_3 = 650 \times 10^6 \text{ hr}^{-1}$$

$$k_4 = 48.2 \text{ hr}^{-1}$$

Next since  $k_3$  is much bigger than  $k_2$  we can approximate the reaction by



Now find  $C_{R\max}$  from Eq 8.8

$$\frac{C_{R\max}}{C_{A0}} = \left( \frac{k_5}{k_6} \right)^{k_6/k_6 - k_5} = \left( \frac{4.8 \times 10^6}{48.2} \right)^{48.2/48.2 - 4.8 \times 10^6} = 0.99988$$

and

$$\bar{t}_{\text{plug}} = \frac{\ln(k_2/k_1)}{k_2 - k_1} = 2.4 \times 10^{-6} \text{ hr} = 0.0086 \text{ s}$$

With such a small time and such a high conversion practically any kind of reactor would be OK but in all cases straight plug flow is best

Do not recycle or bypass

Note: In the article mentioned in the problem statement the minimum residence time allowed was 1.8 s. For this,  $C_R$  is lower than  $C_{R\max}$ . From Eq 8.7 we find, for this time

$$\frac{C_R}{C_{A0}} = \frac{k_5}{k_6 - k_5} \left[ e^{-k_1 \bar{t}} - e^{-k_2 \bar{t}} \right] = (-1) [0 - 0.9762] = 0.9762$$