

a) Overall mass balance:

$$\frac{d(\rho V)}{dt} = w_1 + w_2 - w_3 \tag{1}$$

Energy balance:

$$C \frac{d \left[\rho V(T_3 - T_{ref}) \right]}{dt} = w_1 C(T_1 - T_{ref}) + w_2 C(T_2 - T_{ref}) - w_3 C \left(T_3 - T_{ref}\right)$$
(2)

Because $\rho = \text{constant}$ and $V = \overline{V} = \text{constant}$, Eq. 1 becomes:

$$w_3 = w_1 + w_2 \tag{3}$$

b) From Eq. 2, substituting Eq. 3

$$\rho C \overline{V} \frac{d(T_3 - T_{ref})}{dt} = \rho C \overline{V} \frac{dT_3}{dt} = w_1 C (T_1 - T_{ref}) + w_2 C (T_2 - T_{ref}) - (w_1 + w_2) C (T_3 - T_{ref})$$
(4)

Constants C and T_{ref} can be cancelled:

$$\rho \overline{V} \frac{dT_3}{dt} = w_1 T_1 + w_2 T_2 - (w_1 + w_2) T_3$$
(5)

The simplified model now consists only of Eq. 5.

Solution Manual for Process Dynamics and Control, 4th edition Copyright © 2016 by Dale E. Seborg, Thomas F. Edgar, Duncan A. Mellichamp, and Francis J. Doyle III Degrees of freedom for the simplified model:

Parameters : ρ , \overline{V} Variables : w_1 , w_2 , T_1 , T_2 , T_3 $N_E = 1$ $N_V = 5$

Thus, $N_F = 5 - 1 = 4$

Because w_1 , w_2 , T_1 and T_2 are determined by upstream units, we assume they are known functions of time:

$$w_1 = w_1(t)$$

$$w_2 = w_2(t)$$

$$T_1 = T_1(t)$$

$$T_2 = T_2(t)$$

Thus, N_F is reduced to 0.

2.2

Energy balance:

$$C_{p} \frac{d\left[\rho V(T - T_{ref})\right]}{dt} = wC_{p}(T_{i} - T_{ref}) - wC_{p}(T - T_{ref}) - UA_{s}(T - T_{a}) + Q$$

Simplifying

$$\rho VC_p \frac{dT}{dt} = wC_p T_i - wC_p T - UA_s (T - T_a) + Q$$
$$\rho VC_p \frac{dT}{dt} = wC_p (T_i - T) - UA_s (T - T_a) + Q$$

b) T increases if T_i increases and vice versa.

T decreases if *w* increases and vice versa if $(T_i - T) < 0$. In other words, if $Q > UA_s(T-T_a)$, the contents are heated, and $T > T_i$.

2.3

a) Mass Balances:

$$\rho A_1 \frac{dh_1}{dt} = w_1 - w_2 - w_3 \tag{1}$$

$$\rho A_2 \frac{dh_2}{dt} = w_2 \tag{2}$$

Flow relations:

Let P_1 be the pressure at the bottom of tank 1.

Let P_2 be the pressure at the bottom of tank 2.

Let P_a be the ambient pressure.

Then

$$w_2 = \frac{P_1 - P_2}{R_2} = \frac{\rho g}{g_c R_2} (h_1 - h_2)$$
(3)

$$w_{3} = \frac{P_{1} - P_{a}}{R_{3}} = \frac{\rho g}{g_{c} R_{3}} h_{1}$$
(4)

b) Seven parameters: ρ , A_1 , A_2 , g, g_c , R_2 , R_3

Five variables : h_1 , h_2 , w_1 , w_2 , w_3

Four equations

Thus $N_F = 5 - 4 = 1$

1 input = w_1 (specified function of time) 4 outputs = h_1 , h_2 , w_2 , w_3 Assume constant liquid density, ρ . The mass balance for the tank is

$$\frac{d(\rho Ah + m_g)}{dt} = \rho(q_i - q)$$

Because ρ , A, and m_g are constant, this equation becomes

$$A\frac{dh}{dt} = q_i - q \tag{1}$$

The square-root relationship for flow through the control valve is

$$q = C_v \left(P_g + \frac{\rho g h}{g_c} - P_a \right)^{1/2}$$
(2)

From the ideal gas law,

$$P_g = \frac{(m_g / M)RT}{A(H-h)}$$
(3)

where T is the absolute temperature of the gas.

Equation 1 gives the unsteady-state model upon substitution of q from Eq. 2 and of P_g from Eq. 3:

$$A\frac{dh}{dt} = q_i - C_v \left[\frac{(m_g/M)RT}{A(H-h)} + \frac{\rho gh}{g_c} - P_a\right]^{1/2}$$
(4)

Because the model contains P_a , operation of the system is not independent of P_a . For an open system $P_g = P_a$ and Eq. 2 shows that the system is independent of P_a .

2.5

a)

For linear valve flow characteristics,

$$w_a = \frac{P_d - P_1}{R_a}, \quad w_b = \frac{P_1 - P_2}{R_b}, \quad w_c = \frac{P_2 - P_f}{R_c}$$
 (1)

Mass balances for the surge tanks

$$\frac{dm_1}{dt} = w_a - w_b, \qquad \frac{dm_2}{dt} = w_b - w_c \tag{2}$$

where m_1 and m_2 are the masses of gas in surge tanks 1 and 2, respectively.

If the ideal gas law holds, then

$$P_1 V_1 = \frac{m_1}{M} R T_1, \qquad P_2 V_2 = \frac{m_2}{M} R T_2$$
 (3)

where M is the molecular weight of the gas

 T_1 and T_2 are the temperatures in the surge tanks.

Substituting for m_1 and m_2 from Eq. 3 into Eq. 2, and noticing that V_1 , T_1 , V_2 , and T_2 are constant,

$$\frac{V_1 M}{RT_1} \frac{dP_1}{dt} = w_a - w_b \quad \text{and} \quad \frac{V_2 M}{RT_2} \frac{dP_2}{dt} = w_b - w_c \tag{4}$$

The dynamic model consists of Eqs. 1 and 4.

b) For adiabatic operation, Eq. 3 is replaced by

$$P_1 \left(\frac{V_1}{m_1}\right)^{\gamma} = P_2 \left(\frac{V_2}{m_2}\right)^{\gamma} = C, \text{ a constant}$$
(5)

or

$$m_1 = \left(\frac{P_1 V_1^{\gamma}}{C}\right)^{1/\gamma} \text{ and } m_2 = \left(\frac{P_2 V_2^{\gamma}}{C}\right)^{1/\gamma}$$
(6)

Substituting Eq. 6 into Eq. 2 gives,

$$\frac{1}{\gamma} \left(\frac{V_1^{\gamma}}{C}\right)^{1/\gamma} P_1^{(1-\gamma)/\gamma} \frac{dP_1}{dt} = w_a - w_b$$

$$\frac{1}{\gamma} \left(\frac{V_2^{\gamma}}{C}\right)^{1/\gamma} P_2^{(1-\gamma)/\gamma} \frac{dP_2}{dt} = w_b - w_c$$

as the new dynamic model. If the ideal gas law were not valid, one would use an appropriate equation of state instead of Eq. 3.

2.6

a) Assumptions:

- 1. Each compartment is perfectly mixed.
- 2. ρ and *C* are constant.
- 3. No heat losses to ambient.

<u>Compartment 1</u>: Overall balance (No accumulation of mass):

$$0 = \rho q - \rho q_1 \qquad \text{thus} \qquad q_1 = q \tag{1}$$

Energy balance (No change in volume):

$$V_{1}\rho C \frac{dT_{1}}{dt} = \rho q C (T_{i} - T_{1}) - UA(T_{1} - T_{2})$$
⁽²⁾

Compartment 2:

Overall balance:

$$0 = \rho q_1 - \rho q_2$$
 thus $q_2 = q_1 = q$ (3)

Energy balance:

$$V_2 \rho C \frac{dT_2}{dt} = \rho q C (T_1 - T_2) + U A (T_1 - T_2) - U_c A_c (T_2 - T_c)$$
(4)

b) Eight parameters: ρ , V_1 , V_2 , C, U, A, U_c , A_c Five variables: T_i , T_1 , T_2 , q, T_c Two equations: (2) and (4) Thus $N_F = 5 - 2 = 3$

2 outputs =
$$T_1$$
, T_2
3 inputs = T_i , T_c , q (specify as functions of t)

c) Three new variables: c_i, c_1, c_2 (concentration of species A). Two new equations: Component material balances on each compartment. c_1 and c_2 are new outputs. c_i must be a known function of time.

2.7

As in Section 2.4.2, there are two equations for this system:

$$\frac{dV}{dt} = \frac{1}{\rho} (w_i - w)$$
$$\frac{dT}{dt} = \frac{w_i}{V\rho} (T_i - T) + \frac{Q}{\rho VC}$$

Results:

(a) Since w is determined by hydrostatic forces, we can substitute for this variable in terms of the tank volume as in Section 2.4.5 case 3.

$$\frac{dV}{dt} = \frac{1}{\rho} \left(w_i - C_v \sqrt{\frac{V}{A}} \right)$$
$$\frac{dT}{dt} = \frac{w_i}{\rho V} (T_i - T) + \frac{Q}{\rho VC}$$

This leaves us with the following:

5 variables:
$$V, T, w_i, T_i, Q$$

4 parameters: C, ρ, C_v, A
2 equations

The degrees of freedom are 5-2=3. To make sure the system is specified, we have:

2 output variables: T, V

2 manipulated variables: Q, w_i

1 disturbance variable: T_i

(b) In this part, two controllers have been added to the system. Each controller provides an additional equation. Also, the flow out of the tank is now a manipulated variable being adjusted by the controller. So, we have

4 parameters: C, ρ, T_{sp}, V_{sp} 6 variables: V, T, w_i, T_i, Q, w 4 equations

The degrees of freedom are 6-4=2. To specify the two degrees of freedom, we set the variables as follows:

2 output variables: T, V2 manipulated variables (determined by controller equations): Q, w2 disturbance variables: T_i, w_i

2.8

Additional assumptions:

- (i) Density of the liquid, ρ , and density of the coolant, ρ_J , are constant.
- (ii) Specific heat of the liquid, C, and of the coolant, C_J , are constant.

Because V is constant, the mass balance for the tank is:

$$\rho \frac{dV}{dt} = q_F - q = 0; \text{ thus } q = q_F$$

Energy balance for tank:

$$\rho VC \frac{dT}{dt} = q_F \rho C (T_F - T) - K q_J^{0.8} A (T - T_J)$$
(1)

Energy balance for the jacket:

$$\rho_J V_J C_J \frac{dT_J}{dt} = q_J \rho_J C_J (T_i - T_J) + K q_J^{0.8} A (T - T_J)$$
(2)

where A is the heat transfer area (in ft^2) between the process liquid and the coolant.

Eqs.1 and 2 comprise the dynamic model for the system.

2.9

Assume that the feed contains only A and B, and no C. Component balances for A, B, C over the reactor give.

$$V\frac{dc_{A}}{dt} = q_{i}c_{Ai} - qc_{A} - Vk_{1}e^{-E_{1}/RT}c_{A}$$
(1)

$$V\frac{dc_B}{dt} = q_i c_{Bi} - q c_B + V(k_1 e^{-E_1/RT} c_A - k_2 e^{-E_2/RT} c_B)$$
(2)

$$V\frac{dc_C}{dt} = -qc_C + Vk_2e^{-E_2/RT}c_B$$
(3)

An overall mass balance over the jacket indicates that $q_c = q_{ci}$ because the volume of coolant in jacket and the density of coolant are constant.

Energy balance for the reactor:

$$\frac{d\left[\left(Vc_{A}M_{A}S_{A}+Vc_{B}M_{B}S_{B}+Vc_{C}M_{C}S_{C}\right)T\right]}{dt} = \left(q_{i}c_{Ai}M_{A}S_{A}+q_{i}c_{Bi}M_{B}S_{B}\right)(T_{i}-T) -UA(T-T_{c}) + (-\Delta H_{1})Vk_{1}e^{-E_{1}/RT}c_{A} + (-\Delta H_{2})Vk_{2}e^{-E_{2}/RT}c_{B}$$
(4)

where M_A , M_B , M_C are molecular weights of A, B, and C, respectively

 S_A , S_B , S_C are specific heats of A, B, and C.

U is the overall heat transfer coefficient

A is the surface area of heat transfer

Energy balance for the jacket:

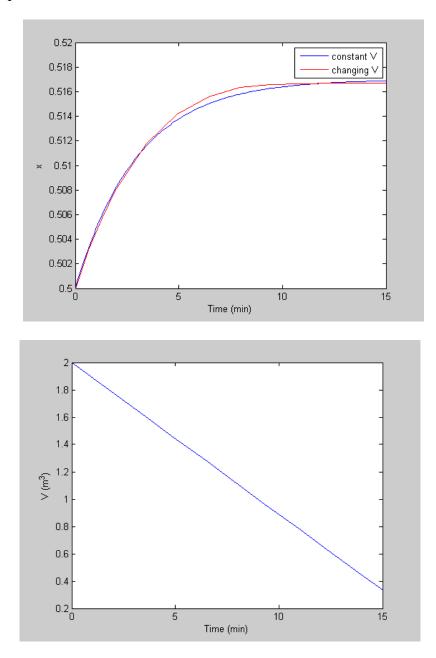
$$\rho_{j}S_{j}V_{j}\frac{dT_{c}}{dt} = \rho_{j}S_{j}q_{ci}(T_{ci} - T_{c}) + UA(T - T_{c})$$
(5)

where:

 ρ_j, S_j are density and specific heat of the coolant. V_j is the volume of coolant in the jacket.

Eqs. 1 - 5 represent the dynamic model for the system.

2.10



The plots should look as shown below:

Notice that the functions are only good for t = 0 to t = 18, at which point the tank is completely drained. The concentration function blows up because the volume function is negative.