

a) Overall mass balance:

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

Energy balance:

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

-w₃C(T₃ - T_{ref}) (2)

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

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

b) From Eq. 2, substituting Eq. 3

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

Constants *C* and *Tref* can be cancelled:

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

The simplified model now consists only of Eq. 5.

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Degrees of freedom for the simplified model:

Parameters : ρ*, 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:

$$
w_1 = w_1(t)
$$

\n
$$
w_2 = w_2(t)
$$

\n
$$
T_1 = T_1(t)
$$

\n
$$
T_2 = T_2(t)
$$

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

$$
\rho V C_p \frac{dT}{dt} = w C_p T_i - w C_p T - U A_s (T - T_a) + Q
$$

$$
\rho V C_p \frac{dT}{dt} = w C_p (T_i - T) - U A_s (T - T_a) + Q
$$

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 $P₁$ 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)

$$
w_3 = \frac{P_1 - P_a}{R_3} = \frac{\rho g}{g_c R_3} h_1 \tag{4}
$$

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

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

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 *Pa*, operation of the system is not independent of *Pa*. 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

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

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,

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

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

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

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.

Compartment 1: 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:

$$
0 = \rho q_1 - \rho q_2 \quad \text{thus} \quad q_2 = q_1 = q \tag{3}
$$

Energy balance:

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

b) Eight parameters: ρ, *V*1, *V*2, *C*, *U*, *A*, *Uc*, *Ac* 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 (specificity 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.

$$
\frac{dV}{dt} = \frac{1}{\rho} \left(w_i - C_v \sqrt{\frac{V}{A}} \right)
$$

$$
\frac{dT}{dt} = \frac{w_i}{\rho V} (T_i - T) + \frac{Q}{\rho V C}
$$

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: *Ti*

(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: $C, \rho, T_{\rm gp}, V_{\rm sp}$ 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 V*, 2 manipulated variables (determined by controller equations): *Q w*, 2 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:

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

Energy balance for tank:

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

Energy balance for the jacket:

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

$$
V\frac{dc_A}{dt} = q_i c_{Ai} - q c_A - V k_1 e^{-E_1/RT} c_A
$$
 (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:

$$
\rho_j S_j V_j \frac{dT_c}{dt} = \rho_j S_j q_{ci} (T_{ci} - T_c) + UA(T - T_c)
$$
\n(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

 0.52 $constant \vee$ 0.518 changing V 0.516 0.514 0.512 0.51 \asymp 0.508 0.506 0.504 0.502 $0.5\frac{V}{0}$ $\overline{5}$ $\overline{10}$ $\frac{1}{15}$ Time (min) $\overline{2}$ 1.8 1.6 1.4 1.2 \times (m³) $0.8\,$ 0.6 0.4 $0.2\frac{L}{0}$ $\overline{5}$ $\overline{10}$ $\overline{15}$ Time (min)

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.