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I certify that the work being submitted is my own, and
that no unauthorized assistance was obtained.

Name (please print): _____

Signature: _____

Exam #1
(Closed book, closed notes, no electronic resources)

1. A continuous vaporizer shown below is a cylindrical pressure vessel of cross-sectional area **A**. The feed is a pure organic liquid with a molecular weight of **M**. The liquid is fed at molar flowrate **N_{in}** and temperature **T_{in}** to the vaporizer that operates at pressure **P** and temperature **T**. Heat added to the vaporizer at a rate of **Q** generates a vapor stream with molar flowrate **N_{out}**. During operation, the liquid holdup of the vessel is **N_L** moles of liquid with a density of **ρ_L**.

- 23/40**
(a) Develop a mathematical model of the system that can be solved to predict the effects of disturbances in liquid feed rate and heat addition rate on the liquid level (**h_L**) in the vaporizer. Assume that the specific heat capacities of the liquid and vapor are both known and ignore the (20 pts) **15**
- (b) Determine how many variables and parameters must be specified to make the model fully solvable. (5 pts) **3**
- (c) On the diagram, draw proposed arrangements for the minimum number of control systems (i.e. sensor/controller + actuator) that would ensure safe operation of this system if significant fluctuations in heating rate and feed rate are expected. Briefly explain your rationale for each control system – i.e. what kind of problem(s) would it prevent. (15 pts) **5**

A = cross sectional area

No reaction

$MW_{\text{avg}} \Rightarrow M$

$$\text{Feed molar flow rate, mol/s} = N_{\text{in}}$$

$$T_{\text{in}} = T \text{ to vaporizer}$$

$$P, T = \text{pressure, } T \text{ of vaporizer}$$

$$\text{Energy balance: } \frac{dQ}{dt} = (FC_p \Delta T)_{\text{in}} - (FC_p \Delta T)_{\text{out}} + Q$$

$$\frac{dQ}{dt} = F_{\text{in}} C_p (T_{\text{in}} - T_{\text{ref}}) - (F_{\text{out}} C_p (T_{\text{out}} - T_{\text{ref}})) + UA(T_2 - T_{\text{bank}}) = T$$

$$\text{Assume } T_{\text{ref}} = 0^\circ\text{C}$$

$$\frac{dQ}{dt} = F_{\text{in}} C_p T_{\text{in}} - F_{\text{out}} C_p T_{\text{out}} + UA(T_2 - T_{\text{ref}}) = MC_p dT/dt$$

$$\text{Mass balance: } \frac{d(N_L)}{dt} = \rho F_{\text{in}} - \rho F_{\text{out}}$$

BACK SIDE

$A = V/h_L$
Quarizer

Sensor ①: concentration sensor - (analyzer)
Senses fluctuations in feed rate ??

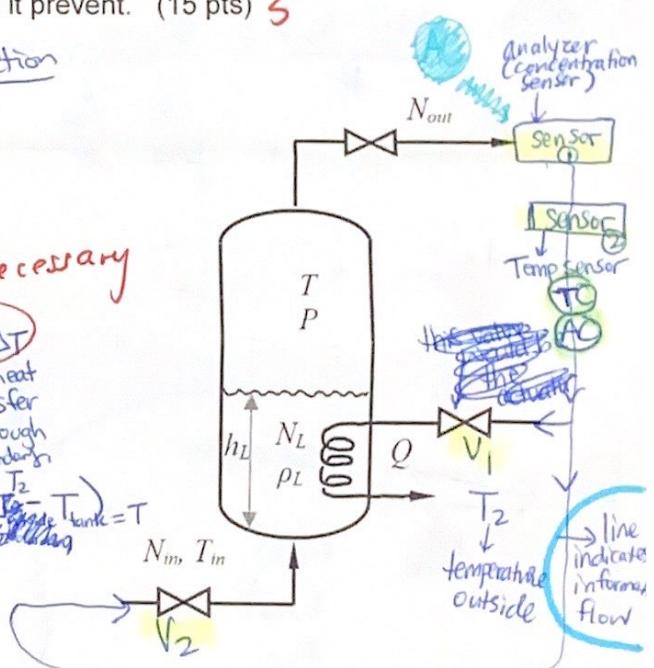
Not clear what measurement used to control what value!

Sensor ②: Temperature sensor - senses heat fluctuation in the outlet stream

Activator: V_1 : valve
Activator: V_2 : By controlling this valve, you can control heat in → by controlling this valve, you can control the heat added to the system to increase or decrease temperature.

By controlling this valve, you can control flow into the system by increasing or decreasing it.

AC: concentration controller (analyzer)



$$\frac{d(P_L \cdot (A \cdot h_L))}{dt} = (P_L \cdot F_{\text{in}}) - (P_L \cdot F_{\text{out}})$$

(assume constant P and A)

$$\frac{d(A \cdot h_L)}{dt} = F_{\text{in}} - F_{\text{out}}$$

(Back side)

The controllers are placed after the sensors to provide the actuators with information about the setpoint.

TC: Temp. controller

Mole balance

(Component balance ~~assumes~~
with feed of pure liquid
being component)

~~dt~~
~~dt~~

(Total mole balance = component mole balance
because of pure feed stream.)

$$\frac{dN_L}{dt} = \frac{d(C \cdot V)}{dt} = \frac{d(C \cdot (A \cdot h_L))}{dt} = \text{In - out } *$$

acc.
of N_L
moles

$$\frac{dN_L}{dt} = \text{In} - \text{Out}$$

$$\frac{dN_L}{dt} = \frac{d(F \cdot C \cdot h_L)}{dt} = F_{in} C_{in} - F_{out} C_{out} = N_{in} - N_{out}$$

(mol/time units) \downarrow \downarrow

F_{in} C_{in} F_{out} C_{out}

mol mol mol mol

not constant

molar flow rates

$$\frac{dN_L}{dt} = N_{in} - N_{out}$$

b) Variables: ~~specified~~ ~~var~~

~~R₂~~

~~N_{in}~~

~~N_{out}~~

~~F_{in}~~

~~C_p~~

~~T_{in}~~

~~T_{out}~~

~~F_{out}~~

~~Q_{heat}~~

Ignore
the
crossed
out variables.

There are
11 unknowns
but C_p is "specified"
So 10 unknowns.

✓

heat eqn continued

$$\frac{dQ}{dt} = \frac{d(MC_p \Delta T)}{dt} = M C_p \frac{dT}{dt}$$

$$\frac{dQ}{dt} = F_{in} C_p T_{in} - F_{out} C_p T_{out} + Q_{heat}$$

When simplified with given variables

11 variables
but 8 are given, so
3 unknown variables
need level sensor
is mass change of
mass balance

eqns
3 balances

- heat/energy balance
- mass balance
- mole balance

1 DOF = $10 - 3 = 7$

The system is
underspecified
and you need
more to solve it

2. BASF Engineering Intern Heinrich Plaumann (no relation to Dr. Heinz Plaumann), was ordered to do a dynamic analysis of outlet temperature in a two-CSTRs-in-series system after a sudden increase in feed concentration. His modeling produced the differential equation below. Mr Plaumann reported to his supervisor that he had analyzed the system using Laplace's Final Value Theorem and had concluded that the reactor temperature would quickly stabilize after the feed disturbance.

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30

$$\frac{d^2T}{dt^2} - 5 \frac{dT}{dt} + 6T = 0 \quad \frac{dT}{dt}(0) = 0, \quad T(0) = 300$$

(a) Find the Laplace transform $T(s)$. (10 pts) 10 $y'(0) = 0 \quad y(0) = 300$

(b) What does the Final Value Theorem predict about temperature as $t \rightarrow \infty$? (10 pts) 10

(c) By comparing $T(s)$ with your transform tables, determine what kinds of time functions should appear in $T(t)$ and discuss Mr. Kastle's prediction. (10 pts) 4

~~for ease of solving~~

$$\frac{d^2T}{dt^2} - 5 \frac{dT}{dt} + 6T = 0$$

$$\times \cancel{T(s)} = F(s)$$

$$\rightarrow \cancel{\times} \cancel{y(s)} = \cancel{I \text{ mean } T(s)} \times$$

Laplace transform of each term:

$$(s^2 F(s) - s f(0) - [\frac{df}{dt}]_{t=0}) - 5 (s F(s) - f(0)) + [6 (F(s))] - \frac{5}{s} = 0$$

$$(s^2 Y(s) - s y(0) - y'(0)) - 5 (s Y(s) - y(0)) + 6 (Y(s)) - \frac{5}{s} = 0 \checkmark$$

$$(s^2 Y(s) - s y(0)) - 5 (s Y(s) - 300) + 6 (Y(s)) - \frac{5}{s} = 0$$

$$s^2 Y(s) - 300s - 5s Y(s) + 1500 + 6Y(s) - \frac{5}{s} = 0$$

~~$Y(s) = T(s)$ used this for ease of solving so I would not miss up~~

$$s^2 (T(s)) - 300s - 5s T(s) + 1500 + 6T(s) \Rightarrow T(s) = \frac{5}{s(s-2)(s-3)} - \frac{1500}{(s-2)(s-3)} + \frac{300s}{(s-2)(s-3)}$$

b) Final value theorem

$$\lim_{t \rightarrow \infty} f(t) = \lim_{s \rightarrow 0} s F(s) \rightarrow \lim_{s \rightarrow 0} \left(\frac{5s}{s(s-2)(s-3)} - \frac{1500s}{(s-2)(s-3)} + \frac{300s^2}{(s-2)(s-3)} \right)$$

$$\Leftrightarrow \frac{5}{(0-2)(0-3)} - 0 + 0 \Rightarrow \frac{5}{(-2)(-3)} = \frac{5}{6}$$

b) Is the final steady state value of the reaction system

$T(t) \rightarrow$ What kind of functions?

$$T(s) = \frac{5}{s(s-2)(s-3)} - \frac{1500}{(s-2)(s-3)} + \frac{300s}{(s-2)(s-3)}$$

\downarrow ① \downarrow ② \downarrow ③

Taking Inverse of each term would give
 $T(t)$ functions such as:

For ① $5 f^{-1} \left\{ \frac{1}{s(s-2)(s-3)} \right\}$

A similar Laplace transform is $n! / (s+a)^{n+1}$ → a constant on top, and if $n=2$ then s will be cubed, since many of the other transforms did not have a constant in the numerator and ~~had~~ an s^3 in the denominator. The inverse is $t^n - at$.

For ② $1500 f^{-1} \left\{ \frac{1}{(s-2)(s-3)} \right\}$

if you take the inverse, it is ~~not~~ similar to the Laplace transform

For ③ $300 f^{-1} \left\{ \frac{s}{(s-2)(s-3)} \right\}$

where a_1, a_2 are -2 and -3
The $f(t)$ is $\frac{1}{a_1 - a_2} (e^{-a_2 t} - e^{-a_1 t})$

A similar Laplace transform is

$\frac{Sta}{(s+a)^2 + \omega^2}$, if $a=0$. There is an s on the top, and s^2 on bottom. The inverse gives

which is an exponential graph because which has an exponential component to the graph

$e^{-at} \cos(\omega t)$, which is exponential and oscillatory

Mr. Plaumann was wrong because the temperature will not quickly stabilize after the disturbance, due to the exponential and oscillatory \downarrow time.

what #s
are a_1, a_2 ? ??

only if quadratic cannot be factored $(s+a_1)(s+a_2)$

Too much overthinking!

$$T(s) = \frac{A}{s} + \frac{B}{s-2} + \frac{C}{s-3}$$

\downarrow
constant \downarrow $e^{2t} \rightarrow \infty$ \downarrow $e^{3t} \rightarrow \infty$