

Engineering 11  
Fall 2003  
James Hunt

Name KEY

Quiz No. 2  
(November 14, 2003)

1 (25 pts) \_\_\_\_\_

2 (25 pts) \_\_\_\_\_

3 (25 pts) \_\_\_\_\_

4 (25 pts) \_\_\_\_\_

mean 62

SD 16

range 94-27

94 - 75 A 14

74 - 63 B 15

62 - 45 C 21

45 - D 8

The exam is closed books and notes.

Do not ask for clarification of the problem statements, part of the exam is understanding the questions.

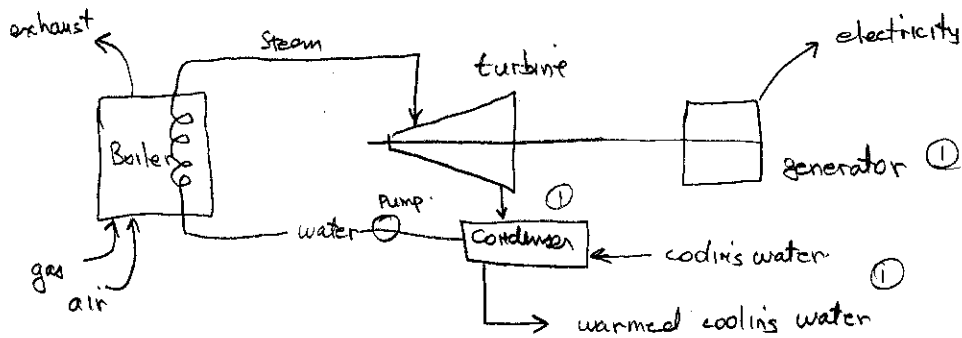
If you think there is an error in the problem, state any necessary assumptions and proceed.

Heat content:  $q = mc_p T$  where  $m$  is mass [kg],  $c_p$  is heat capacity [kJ/kg-K], and  $T$  is temperature [K].

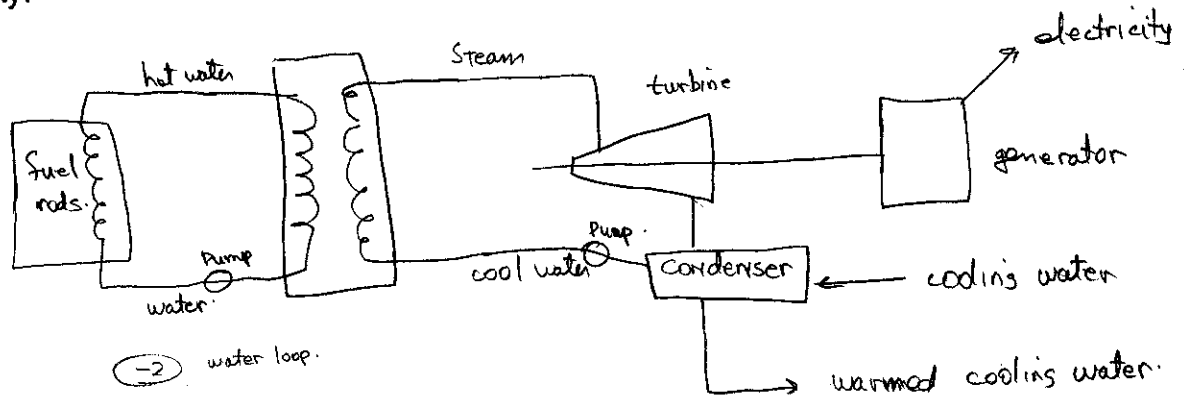
Partial credit is given for partially correct work.

1. Answer the following questions in the space provided.

- (a) Sketch the layout of a natural gas fired electrical generating plant that utilizes steam turbines.



- (b) Sketch the layout of a conventional nuclear power plant used to generate electricity.

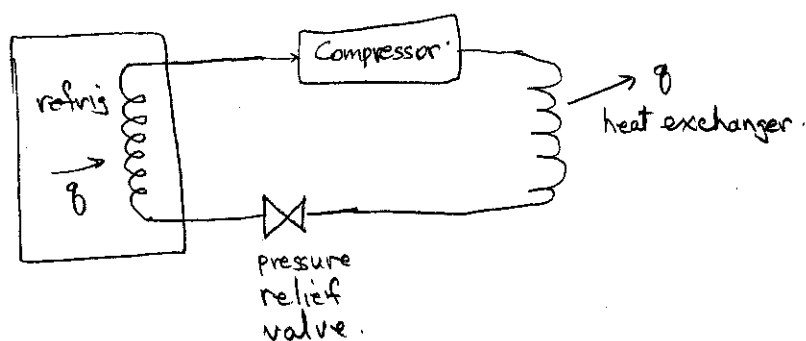


- (c) List **two** reasons why nuclear power plants require more cooling water than fossil-fueled power plants.

- 1) No heat loss out exhaust.
- 2) Operate at lower temperatures, thus less efficient.
- 3) Has two heat transfer steps rather than one for fossil plant.

1. (cont.)

④ (d) Sketch the layout of a refrigerator and describe how heat is removed.



Refrigerant absorbs heat when changes to gas from liquid.

Compressor drives circulation.

External heat exchanger releases heat to environment.

④ (e) How does London smog differ from Los Angeles smog?

London: Smoke + fog. (soot particles, +  $H_2SO_4$  + water drops) (2)

Los Angeles: photochemical smog from  $NO_x$  +  $O_3$  + hydrocarbons. (2)

④ (f) Why is ozone a particularly difficult pollutant to control in urban air basins?

There is no direct source of ozone, it is a secondary pollutant formed from the interaction of  $NO$ , hydrocarbons and sunlight.

Thus need control of  $NO_x$  and hydrocarbons that could have separate sources.

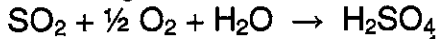
$O_3$  is difficult to model, thus difficult to set regulations.

2. Urban air quality is impacted by  $\text{SO}_2$  in many locations.

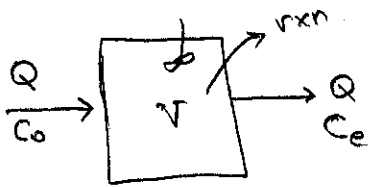
⑤ (a) What is the predominant source of  $\text{SO}_2$  found in air?

combustion of coal.

⑤ (b)  $\text{SO}_2$  undergoes first order oxidation to sulfuric acid:



with a first order rate constant,  $k_s$  [ $\text{s}^{-1}$ ]. Air flows into a well-mixed valley with a flow rate of  $Q$  [ $\text{m}^3/\text{s}$ ] and the air volume in the valley is  $V$  [ $\text{m}^3$ ]. The  $\text{SO}_2$  concentration entering the valley is  $C_0$  [ $\text{mol}/\text{m}^3$ ] and there is no other source of  $\text{SO}_2$  within the valley. Derive an expression for the  $\text{SO}_2$  concentration leaving the valley under steady state conditions.



$\text{SO}_2$  balance:

moles in = moles out + moles removed by rxn.  
by flow by flow

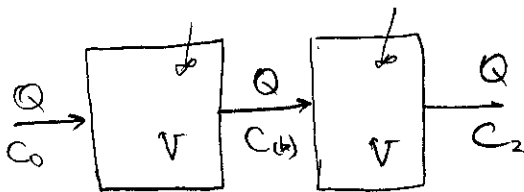
$$Q C_0 \Delta t = Q C_e \Delta t + k_s C_e V \Delta t.$$

$$C_e = \frac{Q C_0}{Q + k_s V}$$

$$= \frac{C_0}{1 + k_s \frac{V}{Q}} = \frac{C_0}{1 + k_s \tau}$$

$C_e \neq C_{\text{inside}}$  ②

⑤ (c) Downwind of the valley described in part (b), there is another identical valley of volume  $V$ . What is the expression for the  $\text{SO}_2$  concentration leaving the second valley under steady state conditions?



repeat process in (b)

$$C_2 = \frac{C_1}{1 + k_s \tau}$$

$$= \frac{C_0}{(1 + k_s \tau)^2}$$

2. (cont.)

- ④ (d) What is the concentration of  $\text{H}_2\text{SO}_4$  leaving the second valley described in part (c)? There is no deposition of  $\text{H}_2\text{SO}_4$  to the land surface in either valley.

$$\text{moles } (\text{H}_2\text{SO}_4) + \text{moles } \text{SO}_2 = \text{moles } \text{SO}_2 \text{ entering system.} \quad \textcircled{2}$$

$$\begin{aligned} C_{\text{H}_2\text{SO}_4} &= C_0 - C_2 \\ &= C_0 \left( 1 - \frac{1}{(1 + k_s \tau)^2} \right) \end{aligned} \quad \textcircled{3}$$

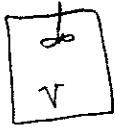
- ⑤ (e) How would an increase in air flow rate through the valleys change your answers in (b) and (c)?

IF  $Q$  increase,  $\tau$  decreases and  $C$  increases, less conversion because less time to react.

3. Nuclear power plants generate radioactive wastes that are fission products that undergo decay and release heat. This process is represented as a first order reaction  
 $A \rightarrow B$

Where radioisotope A decays to a stable isotope B with a first order rate constant  $k_A$ . The reaction generates heat,  $H$  [kJ/mol]. You have a closed and thermally insulated vessel containing A at an initial concentration of  $C_{A,0}$  [mol/m<sup>3</sup>] dissolved in water.

- ⑧ (a) Derive an expression for the concentration of A within the vessel.



in = out + accumulated.

$$0 = k_A C_A V \Delta t + \Delta C_A V \quad \text{④}$$

$$\frac{\Delta C_A}{\Delta t} = -k_A C_A$$

$$\frac{dC_A}{dt} = -k_A C_A \quad \downarrow \text{ok.}$$

$$\text{integrate with i.c. } C_A(t=0) = C_{A,0} \quad \text{④}$$

$$C_A(t) = C_{A,0} e^{-k_A t} \quad \text{② only}$$

- ⑧ (b) Derive an expression for the concentration of B within the vessel.

By mole balance.

$$C_B(t) = C_{A,0} - C_A(t)$$

$$C_B(t) = C_{A,0} (1 - e^{-k_A t})$$

- ⑧ (c) If the vessel had an initial temperature of  $T_0$ , what is the expression for the steady state temperature?

heat balance.

initial heat + heat added by decay = final heat

$$m C_p T_0 + H \cdot C_{A,0} \cdot V = m C_p T_f$$

$$\left[ \frac{\text{kJ}}{\text{mole}} \right] \left( \frac{\text{mole}}{\text{m}^3} \right) (\text{m}^3)$$

$$T_f = \frac{m C_p T_0 + H C_{A,0} V}{m C_p}$$

$$T_f = T_0 + \frac{H C_{A,0} V}{m C_p}$$

$$T_f = T_0 + \frac{H C_{A,0}}{\rho C_p}$$

$$\rho = \frac{m}{V}$$