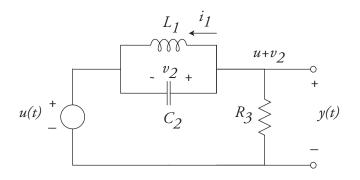
#### Unified Engineering I

## Problem S13 (Signals and Systems) SOLUTION

It's easiest to use the node method for this problem. Label the states  $i_1$ , and  $v_2$ , and the nodes as below:



The state vector is defined as

$$\underline{x} = \left[ \begin{array}{c} i_1 \\ v_2 \end{array} \right]$$

So we need to find  $di_1/dt$ ,  $dv_2/dt$ . Start with the inductor, with

$$\frac{di_1}{dt} = \frac{v_1}{L_1} = \frac{v_2}{L_1}$$

since  $v_2 = v_1$ . For the capacitor,

$$\frac{dv_2}{dt} = \frac{i_2}{C_2}$$

To find  $i_2$ , apply KCL at the  $u + v_2$  node:

$$i_1 + i_2 + \frac{u + v_2}{R_3} = 0$$

Solving for  $i_2$ ,

$$i_2 = -i_1 - \frac{u + v_2}{R_3}$$

Therefore,

$$\frac{dv_2}{dt} = -\frac{i_1}{C_2} - \frac{u + v_2}{R_3 C_2}$$

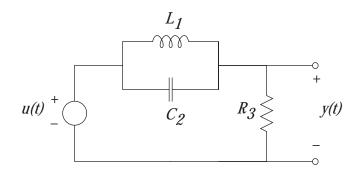
In state-space form,

$$\frac{d}{dt} \begin{bmatrix} i_1 \\ v_2 \end{bmatrix} = \begin{bmatrix} 0 & 1/L_1 \\ -1/C_2 & -1/R_3C_2 \end{bmatrix} \begin{bmatrix} i_1 \\ v_2 \end{bmatrix} + \begin{bmatrix} 0 \\ -1/R_3C_2 \end{bmatrix} u$$
$$y = \begin{bmatrix} 0 & 1 \end{bmatrix} \begin{bmatrix} i_1 \\ v_2 \end{bmatrix} + \begin{bmatrix} 1 \end{bmatrix} u$$

## Unified Engineering I

# Problem S14 (Signals and Systems) SOLUTION

Consider the RLC circuit of Problem S13, shown below:



1. From S13, we found a state-space description to be

$$\frac{d}{dt} \begin{bmatrix} i_1 \\ v_2 \end{bmatrix} = \begin{bmatrix} 0 & 1/L_1 \\ -1/C_2 & -1/R_3C_2 \end{bmatrix} \begin{bmatrix} i_1 \\ v_2 \end{bmatrix} + \begin{bmatrix} 0 \\ -1/R_3C_2 \end{bmatrix} u$$
$$y = \begin{bmatrix} 0 & 1 \end{bmatrix} \begin{bmatrix} i_1 \\ v_2 \end{bmatrix} + \begin{bmatrix} 1 \end{bmatrix} u$$

Therefore,

$$sI - A = \begin{bmatrix} s & -1/L_1 \\ 1/C_2 & s + 1/R_3C_2 \end{bmatrix}$$

Inverting,

$$(sI - A)^{-1} = \frac{1}{s^2 + s/R_3C_2 + 1/L_1C_2} \begin{bmatrix} s + 1/R_3C_2 & +1/L_1 \\ -1/C_2 & s \end{bmatrix}$$

Multiplying by C,

$$C(sI - A)^{-1} = \frac{1}{s^2 + s/R_3C_2 + 1/L_1C_2} \begin{bmatrix} -1/C_2 & s \end{bmatrix}$$

Multiplying by B,

$$C(sI - A)^{-1}B = \frac{-s/R_3C_2}{s^2 + s/R_3C_2 + 1/L_1C_2}$$

Adding D,

$$G(s) = C(sI - A)^{-1}B + D = \frac{-s/R_3C_2}{s^2 + s/R_3C_2 + 1/L_1C_2} + 1 = \frac{s^2 + 1/L_1C_2}{s^2 + s/R_3C_2 + 1/L_1C_2}$$

2. To find the transfer function using impedance methods, note that the inductor and capacitor have parallel impedance

$$Z_{LC} = Z_L ||Z_C = \frac{Z_L Z_C}{Z_L + Z_C} = \frac{Ls \left(\frac{1}{C_s}\right)}{Ls + \frac{1}{C_s}} = \frac{Ls}{LCs^2 + 1}$$

(Note that I have dropped the numerical subscripts, since no confusion can arise.) Then the circuit is a voltage divider, with transfer function

$$G(s) = \frac{R}{Z_{LC} + R} = \frac{R}{\frac{Ls}{LCs^2 + 1} + R} = \frac{R(LCs^2 + 1)}{R(LCs^2 + 1) + Ls} = \frac{RLCs^2 + R}{RLCs^2 + RLs + R}$$

Dividing numerator and denominator by RLC gives

$$G(s) = \frac{s^2 + 1/L_1C_2}{s^2 + s/R_3C_2 + 1/L_1C_2}$$

as in part (1).

3. For component values

$$L_1 = 1$$
 H,  $C_2 = 0.25$  F,  $R_3 = 10 \Omega$ 

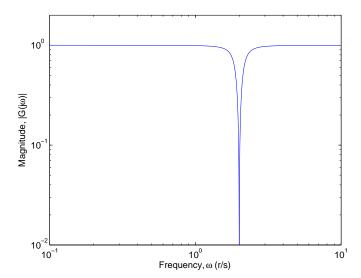
the transfer function is

$$G(s) = \frac{s^2 + 4}{s^2 + 0.4s + 4}$$

Therefore,

$$G(j\omega) = \frac{-\omega^2 + 4}{-\omega^2 + 0.4j\omega + 4}$$

You are then to plot the magnitude of this function. This could be done in a spread sheet, by calculating the transfer function frequency by frequency, or by using the Matlab command **bode**. I used Matlab. (Most of you will probably use a spreadsheet, since we haven't used Matlab this term.) My plot is shown below:



Note that I've plotted the magnitude on a log-log scale. (This is how experts always plot transfer functions.) Most of you will probably plot on a linear-linear scale — that's OK. On either scale, it's clear why this is a "notch filter" — graphically, there's a notch in the plot which is otherwise a straight line.

#### Problem S15 (Signals and Systems) SOLUTION

In Problem S14, we found that

$$G(s) = \frac{s^2 + 1/L_1C_2}{s^2 + s/R_3C_2 + 1/L_1C_2}$$

For component values

$$L_1 = 1$$
 H,  $C_2 = 0.25$  F,  $R_3 = 10 \ \Omega$ 

the transfer function is

$$G(s) = \frac{s^2 + 4}{s^2 + 0.4s + 4}$$

1. For the input

$$u(t) = \cos t$$

the complex frequency is  $s = j\omega$ , where  $\omega = 1$ , and the complex amplitude is U = 1. Therefore,

$$Y = G(s)U = G(1j) \cdot 1 \approx 0.9825 - 0.1310j$$

The real and imaginary parts of the amplitude correspond to the cosine and (minus) sine amplitudes. Therefore,

$$u(t) \approx 0.9825 \cos t + 0.1310 \sin t$$

2. For the input

$$u(t) = 3\sin 2t$$

the complex frequency is  $s = j\omega$ , where  $\omega = 2$ , and the complex amplitude is U = 3. Therefore,

$$Y = G(s)U = G(2j) \cdot 3 = 0$$

The amplitude of y is identically zero, because the frequency of the input signal is at exactly the notch frequency. Therefore,

$$u(t) = 0$$

3. For the input

$$u(t) = 2\cos 4t + \sin 4t$$

the complex frequency is  $s = j\omega$ , where  $\omega = 4$ , and the complex amplitude is U = 2 - j. Therefore,

$$Y = G(s)U = G(4j) \cdot (2-j) \approx (0.9825 + 0.1310j)(2-j) \approx 2.0961 - 0.7205j$$

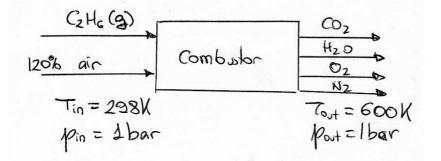
Finally,

$$u(t) \approx 2.0961 \cos 4t + 0.7205 \sin 4t$$

Fall 2006

# Problem 4

GIVEN:



**ASSUMPTIONS:** • Steady flow.

• Neglect kinetic and potential energy effects.

#### CONCEPTS: •

- Stoichiometry.First Law in CV form.
- Enthalpy of formation.

### SOLUTION:

a) The stoichiometric reaction is:

$$C_2H_6 + 3.5(O_2 + 3.76N_2) \rightarrow 2CO_2 + 3H_2O + 3.5 \times 3.76N_2$$

With 20% excess air:

$$C_2H_6 + 4.2(O_2 + 3.76N_2) \rightarrow 2CO_2 + 3H_2O + 0.7O_2 + 4.2 \times 3.76N_2$$

Then, the air/fuel ratio is:

$$\frac{1}{f} = \frac{m_{\rm air}}{m_{\rm fuel}} = \frac{4.2(M_{\rm O_2} + 3.76M_{\rm N_2})}{M_{\rm C_2H_6}} = 19.22$$

b) Per kmol of fuel, the products will be:

- 2 kmol of  $CO_2$ .
- 3 kmol of  $H_2O$ .
- 0.7 kmol of  $O_2$ .
- 15.79 kmol of N<sub>2</sub>.

c) The enthalpy of a reaction at the standard conditions is:

$$\begin{split} \Delta \overline{h}_{\text{reaction}} &= 2\overline{h}_{f,\text{CO}_2}^0 + 3\overline{h}_{f,\text{H}_2\text{O}}^0 + 0.7\overline{h}_{f,\text{O}_2}^0 + 15.79\overline{h}_{f,\text{N}_2}^0 - \\ &- \overline{h}_{f,\text{C}_2\text{H}_6}^0 - 4.2\overline{h}_{f,\text{O}_2}^0 - 15.79\overline{h}_{f,\text{N}_2}^0 = \\ &= 2\overline{h}_{f,\text{CO}_2}^0 + 3\overline{h}_{f,\text{H}_2\text{O}}^0 - \overline{h}_{f,\text{C}_2\text{H}_6}^0 = -1.428 \times 10^6 \text{ kJ/kmol} \end{split}$$

d) The amount of heat transfer per kmol of fuel is obtained from the first law:

$$\overline{q} = \sum_{R} n_R \overline{h}_R - \sum_{P} n_P \overline{h}_P$$

The enthalpy of the reactants is:

$$\sum_{R} n_{R}\overline{h}_{R} = \overline{h}_{C_{2}H_{6}} + 4.2\overline{h}_{O_{2}} + 15.79\overline{h}_{N_{2}} = \overline{h}_{f,C_{2}H_{6}}^{0} + 4.2\overline{h}_{f,O_{2}}^{0} + 15.79\overline{h}_{f,N_{2}}^{0} = \overline{h}_{f,C_{2}H_{6}}^{0}$$

The enthalpy of the products is:

$$\sum_{P} n_{P} \overline{h}_{P} = 2\overline{h}_{CO_{2}} + 3\overline{h}_{H_{2}O} + 0.7\overline{h}_{O_{2}} + 15.79\overline{h}_{N_{2}},$$

where

$$\overline{h}_{\rm CO_2} = \overline{h}_{f,\rm CO_2}^0 + \Delta \overline{h}_{\rm CO_2}|_{T=600 \text{ K}}$$

$$\overline{h}_{\rm H_2O} = \overline{h}_{f,\rm H_2O}^0 + \Delta \overline{h}_{\rm H_2O}|_{T=600 \text{ K}}$$

$$\overline{h}_{\rm O_2} = \overline{h}_{f,\rm O_2}^0 + \Delta \overline{h}_{\rm O_2}|_{T=600 \text{ K}} = \Delta \overline{h}_{\rm O_2}|_{T=600 \text{ K}}$$

$$\overline{h}_{\rm N_2} = \overline{h}_{f,\rm N_2}^0 + \Delta \overline{h}_{\rm N_2}|_{T=600 \text{ K}} = \Delta \overline{h}_{\rm N_2}|_{T=600 \text{ K}}$$

Then:

$$\sum_{P} n_{P} \overline{h}_{P} = (2\overline{h}_{f,CO_{2}}^{0} + 3\overline{h}_{f,H_{2}O}^{0}) + (2\Delta\overline{h}_{CO_{2}} + 3\Delta\overline{h}_{H_{2}O}^{0} + 0.7\Delta\overline{h}_{O_{2}} + 15.79\Delta\overline{h}_{N_{2}})|_{T=600 \text{ K}}$$

 $\Delta \overline{h}_{CO_2}$ ,  $\Delta \overline{h}_{H_2O}$ ,  $\Delta \overline{h}_{O_2}$  and  $\Delta \overline{h}_{N_2}$  are the enthalpy change between the reference state and the substance at the exit conditions ( $T_{out} = 600 \text{ K}$ ,  $p_{out} = 1 \text{ bar}$ ). These  $\Delta \overline{h}$  can be found in tables:

$$\begin{split} \Delta h_{\rm CO_2} &= 12906 \text{ kJ/kmol} \\ \Delta \overline{h}_{\rm H_2O} &= 10499 \text{ kJ/kmol} \\ \Delta \overline{h}_{\rm O_2} &= 9245 \text{ kJ/kmol} \\ \Delta \overline{h}_{\rm N_2} &= 8894 \text{ kJ/kmol} \end{split}$$

Adding the different enthalpy of the different products and reactants, and using the definition of  $\Delta \overline{h}_{\text{reaction}}$ :

$$\overline{q} = -\Delta \overline{h}_{\text{reaction}} - 2\Delta \overline{h}_{\text{CO}_2} - 3\Delta \overline{h}_{\text{H}_2\text{O}}^0 - 0.7\Delta \overline{h}_{\text{O}_2} - 15.79\Delta \overline{h}_{\text{N}_2} = 1.224 \times 10^6 \text{ kJ/kmol}$$

The solutions by WAITE  

$$\frac{Tre 3 cock}{Pe 95 x 10^{3} Pa}$$

$$\frac{Tre 4 cock}{Pe 95 x 10^$$

$$T_{T} = \text{const} \text{ if a diabatic } \text{ is no external work}$$
  
So 
$$T_{T_{out}} = 300 \text{ K}$$
  

$$T_{\text{Tout}} = T_{\text{out}} + \frac{C^{2}}{2C_{p}} \implies C_{\text{out}} = 118 \frac{\text{M}}{\text{s}}$$
  

$$\left(\frac{T_{\text{Tout}}}{T_{\text{out}}}\right) = \frac{P_{T_{\text{out}}}}{P_{\text{out}}} \left(\frac{300}{293.06}\right)^{T/s-1} \cdot 100 \times 10^{3} = 108546 \text{ Pa}$$
  

$$\left(\frac{P_{\text{Tout}}}{P_{\text{out}}}\right) = \frac{P_{T_{\text{out}}}}{P_{\text{out}}} \left(\frac{300}{293.06}\right)^{T/s-1} \cdot 100 \times 10^{3} = 108546 \text{ Pa}$$

TI3 SOLUTIONS BY WAITZ

a)  $S_2 - S_1 = C_V \ln\left(\frac{T_2}{T_1}\right) + R \ln\left(\frac{V_2}{V_1}\right) = 10085 \ln\left(\frac{270}{300}\right) + 4124 \ln\left(\frac{200}{100}\right)$ b) FOR QUASI-STATIC ADIABATIC  $\underline{T}_{2} = \left(\frac{V_{1}}{V_{2}}\right)^{r-1} = \left(\frac{100}{200}\right)^{r} = (0.753)$ Tz = 226 K W = -10085 (226-300) = 746290 J/kg As=0 C) IRREVERSIBLE PROCESS: T,= 300K, T== 288K V1=0.0861 kg , V2=0.1722 kg AS = Cvln (F) + Rln (V2) = 716.5 ln (300) + 287 ln (0.1722) 45 irrev. = 169.7J REVENSIBLE PROCESS: T. = 300K Tz = 227.44 VI = 0.0861 Mg Vz = 0.1722 M3/kg AS = 716.50 (227.4) + 287 ln (0.1722) = 0 V