

Multiple Choice Sections:

- ① For a reversible & adiabatic process, $\Delta S = 0$
 \hookrightarrow (b) S_e always equals S_i

- ② $m = 1 \text{ kg}$
 $T_2 = T_1$
 $P_2 = 2P_1$
- Assumptions:
 • Const specific heats

$$S_2 - S_1 = C_p \ln \left| \frac{T_2}{T_1} \right| - R \ln \left| \frac{P_2}{P_1} \right|$$

$$= 0 - R \ln |2| \rightarrow \boxed{S_2 - S_1 = -R \ln(2)}$$

- ③ $v = 0.001 \text{ m}^3/\text{kg}$
 $P_1 = 100 \text{ kPa}$
 $P_2 = 300 \text{ kPa}$

Assumptions:

- $\Delta KE = 0$
- $\Delta PE = 0$
- v is constant through a pump

Min Work = Reversible work: $W_{rev} = \int_1^2 v dp$

$$W_{rev} = v_1 \int_1^2 dp = v_1 (P_2 - P_1) = 0.001 (300 - 100)$$

$$\boxed{W_{rev} = 0.200 \text{ kJ/kg}}$$

- ④ A Carnot cycle may not contain process that are irreversible

- ⑤ $T_H = 20^\circ\text{C}$
 $T_L = 0^\circ\text{C}$

$$\eta_{\text{carnot}} = 1 - \frac{T_L}{T_H} = 1 - \frac{(0+273)}{(20+273)}$$

$$\boxed{\eta_c = 1 - \frac{(273)}{293}}$$

Free Response:

① Given:

$$\begin{aligned} m &= 0.5 \text{ kg} \\ p_1 &= 100 \text{ kPa} \\ T_1 &= 10^\circ\text{C} \\ R &= 2.077 \text{ kJ/kg}\cdot\text{K} \quad (\text{ideal gas}) \\ c_v &= 3.116 \text{ kJ/kg}\cdot\text{K} \quad (\text{constant specific heats}) \\ p_0 &= 500 \text{ kPa} = \text{atmospheric pressure} \end{aligned}$$

Assumptions

- Ideal gas w/ const c_p/c_v

a) Piston will begin to move when the pressure in the piston exceeds the atmospheric pressure

$$\hookrightarrow p_2 = 500 \text{ kPa}$$

$$\hookrightarrow v_2 = v_1 \quad (\text{NO volume change before piston moves})$$

I Law for a closed system:

$$Q_{12} = \Delta U_{12} + W_{\text{exp}}$$

- neglect expansion work since volume doesn't change.

$$Q_{12} = \Delta U_{12} = m c_v \Delta T_{12}$$

- using equation for const specific heat

$$p_1 v_1 = R T_1 \rightarrow \frac{p_1}{T_1} = \text{const} \rightarrow \frac{p_1}{T_1} = \frac{p_2}{T_2} \quad \bullet \text{ use of ideal gas law}$$

$$\hookrightarrow \frac{100 \text{ kPa}}{(10+273)} = \frac{500 \text{ kPa}}{T_2} \rightarrow T_2 = 1415 \text{ K} = 1142^\circ\text{C}$$

$$\rightarrow Q_{12} = \Delta U_{12} = m c_v \Delta T_{12} = (0.5)(3.116)(1142 - 10)$$

$$\boxed{Q_{12} = 1763.656 \text{ kJ}}$$

$$b) \Delta S_{12} = m(S_2 - S_1) = m \left[c_v \ln \left| \frac{T_2}{T_1} \right| + R \ln \left| \frac{v_2}{v_1} \right| \right] \quad \text{(Constant volume)}$$

- Entropy change for ideal gas w/ const specific heats.

$$= (0.5) \left[3.116 \ln \left| \frac{1415}{293} \right| \right]$$

$$\boxed{\Delta S_{12} = 2.507 \frac{\text{kJ}}{\text{K}}}$$

$$\begin{aligned} c) \Delta S_{\text{res}} &= \frac{Q_{\text{res}}}{T_R} = -\frac{Q_{12}}{T_R} \\ &= \frac{-1763.56}{2000} = \boxed{-0.882 \frac{\text{kJ}}{\text{K}}} \end{aligned}$$

- NO, they should not be equal. A heat transfer over a finite ΔT is an irreversible process and will lead to some entropy generation.

② Given:

- adiabatic turbine ($\dot{Q} = 0$)
 - Steady state (Neglect ΔKE and ΔPE)
 - $P_1 = 3.0 \text{ MPa}$
 - $T_1 = 300^\circ\text{C}$
 - $\dot{m}_1 = 14.0 \text{ kg/s}$
 - $P_2 = 0.5 \text{ MPa}$ at sat. vapor state
 - $P_3 = 5.0 \text{ kPa}$
 - Process $1 \rightarrow 2$ is irreversible
 - Process $2 \rightarrow 3$ is reversible
- } Assumptions

a) State 1: At 300°C , $P_{\text{sat}} = 8.547 \text{ MPa}$ (Table A-4)
 $\hookrightarrow P_1 < P_{\text{sat}}$ so this is a superheated vapor

$$\rightarrow \text{at } 3 \text{ MPa, } 300^\circ\text{C: } h_1 = 2994.3 \text{ kJ/kg (Table A-6)}$$
$$s_1 = 6.5412 \text{ kJ/kg}\cdot\text{k}$$

State 2: At 0.5 MPa , sat. vapor state:

$$h_2 = h_g(0.5 \text{ MPa}) = 2748.1 \text{ kJ/kg (Table A-5)}$$
$$s_2 = s_g(0.5 \text{ MPa}) = 6.8207 \text{ kJ/kg}\cdot\text{k}$$

State 3: Given that process $2-3$ is reversible and adiabatic, this is an isentropic process:

$$s_3 = s_2 = 6.8207 \text{ kJ/kg}\cdot\text{k}$$

$$\hookrightarrow \text{At } P_3 = 5.0 \text{ kPa, } s_3 = 6.8207 \text{ kJ/kg}\cdot\text{k} =$$

$$\left. \begin{array}{l} s_f = 0.4762 \\ s_g = 8.3938 \end{array} \right\} s_f < s < s_g \rightarrow \text{2-phase mixture}$$

(Table A-5)

$$\text{- Find quality: } x = \frac{s_3 - s_f}{s_g - s_f} = \frac{6.8207 - 0.4762}{8.3938 - 0.4762} = 0.801$$

- Use quality to determine h_3 :

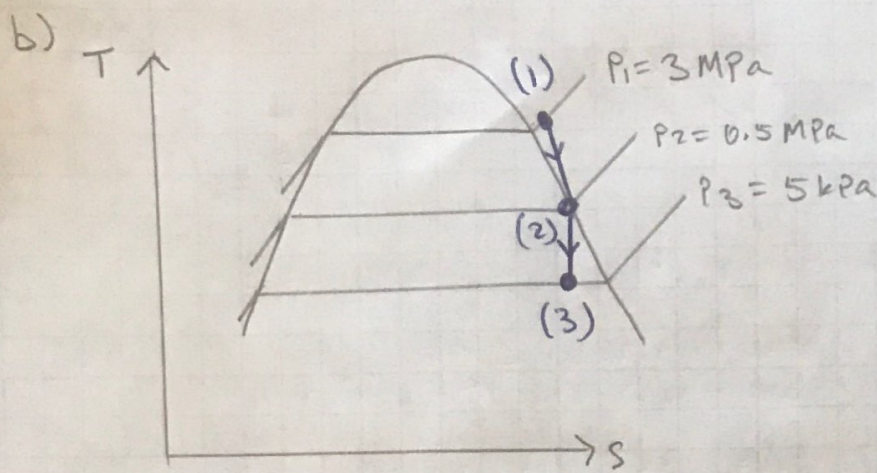
$$h_3 = h_f + x(h_{fg}) \rightarrow h_f = 137.75 \text{ kJ/kg}$$
$$h_{fg} = 2423.0 \text{ kJ/kg}$$

$$\hookrightarrow h_3 = 137.75 + (0.801)(2423.0)$$

$$h_3 = 2079.34 \text{ kJ/kg}$$
$$s_3 = 6.8207 \text{ kJ/kg}\cdot\text{k}$$

$$h_1 = 2994.3 \text{ kJ/kg}$$
$$s_1 = 6.5412 \text{ kJ/kg}\cdot\text{k}$$

$$h_2 = 2748.1 \text{ kJ/kg}$$
$$s_2 = 6.8207 \text{ kJ/kg}\cdot\text{k}$$



c) Set up open system balance equations: steady state

I Law balance: $\dot{Q}_{in} + \dot{W}_{in} + \sum \dot{m}_i (h_i + KE_i + PE_i)$
 $= \dot{Q}_{out} + \dot{W}_{out} + \sum \dot{m}_e (h_e + KE_e + PE_e)$

• neglect \dot{W}_{in} , \dot{Q}_{in} , \dot{Q}_{out} , KE , PE

$\dot{m}_1 h_1 = \dot{W}_{out} + \dot{m}_2 h_2 + \dot{m}_3 h_3$ Given: $\dot{W}_{out} = 11800 \text{ kW}$

Mass Balance: $\sum \dot{m}_{in} = \sum \dot{m}_{out} \rightarrow \dot{m}_1 = \dot{m}_2 + \dot{m}_3$
 • Conservation of mass $14 = \dot{m}_2 + \dot{m}_3 \rightarrow 14 - \dot{m}_2 = \dot{m}_3$

$\rightarrow (14)(2994.3) = 11800 + \dot{m}_2(2748.1) + (14 - \dot{m}_2)(2079.4)$

$= 11800 + \dot{m}_2(2748.1) + 14(2079.4) - \dot{m}_2(2079.4)$

$1008.64 = \dot{m}_2(668.7) \rightarrow \dot{m}_2 = 1.508 \text{ kg/s}$
 $\dot{m}_3 = 12.49 \text{ kg/s}$

d) II Law Open System balance:

$\dot{S}_{gen} = \sum \dot{m}_e s_e - \sum \dot{m}_i s_i - \frac{\dot{Q}}{T} \rightarrow 0 \text{ (adiabatic)}$
 $= \dot{m}_2 s_2 + \dot{m}_3 s_3 - \dot{m}_1 s_1$
 $= 1.508(6.8207) + 12.49(6.8207) - 14(6.5412)$

$\dot{S}_{gen} = 3.913 \frac{\text{kW}}{\text{K}}$