



بارم نمره هر سوال ۲/۸۰ می باشد.

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$$h_1 = h_f @ 10 \text{ kPa} = 191.81 \text{ kJ/kg}$$

$$v_1 = v_f @ 10 \text{ kPa} = 0.00101 \text{ m}^3/\text{kg}$$

$$w_{pl,in} = v_1 (P_2 - P_1) \\ = (0.00101 \text{ m}^3/\text{kg})(600 - 10 \text{ kPa}) \left(\frac{1 \text{ kJ}}{1 \text{ kPa} \cdot \text{m}^3} \right) \\ = 0.60 \text{ kJ/kg}$$

$$h_2 = h_1 + w_{pl,in} = 191.81 + 0.60 = 192.41 \text{ kJ/kg}$$

$$h_3 = h_f @ 0.6 \text{ MPa} = 670.38 \text{ kJ/kg}$$

Mixing chamber:

$$\dot{E}_{in} - \dot{E}_{out} = \Delta \dot{E}_{system} \stackrel{\text{steady}}{\approx} 0 \rightarrow \dot{E}_{in} = \dot{E}_{out}$$

$$\sum \dot{m}_i h_i = \sum \dot{m}_e h_e \rightarrow \dot{m}_4 h_4 = \dot{m}_2 h_2 + \dot{m}_3 h_3$$

$$\text{or, } h_4 = \frac{\dot{m}_2 h_2 + \dot{m}_3 h_3}{\dot{m}_4} = \frac{(22.50)(192.41) + (7.50)(670.38)}{30} = 311.90 \text{ kJ/kg}$$

$$v_4 \approx v_f @ h_4 = 311.90 \text{ kJ/kg} = 0.001026 \text{ m}^3/\text{kg}$$

$$w_{pll,in} = v_4 (P_5 - P_4) \\ = (0.001026 \text{ m}^3/\text{kg})(7000 - 600 \text{ kPa}) \left(\frac{1 \text{ kJ}}{1 \text{ kPa} \cdot \text{m}^3} \right) \\ = 6.57 \text{ kJ/kg}$$

$$h_5 = h_4 + w_{pll,in} = 311.90 + 6.57 = 318.47 \text{ kJ/kg}$$

$$P_6 = 7 \text{ MPa} \left. \begin{array}{l} h_6 = 3411.4 \text{ kJ/kg} \\ T_6 = 500^\circ\text{C} \end{array} \right\} s_6 = 6.8000 \text{ kJ/kg} \cdot \text{K}$$

$$P_7 = 0.6 \text{ MPa} \left. \begin{array}{l} h_7 = 2774.6 \text{ kJ/kg} \\ s_7 = s_6 \end{array} \right\}$$

$$P_8 = 10 \text{ kPa} \left. \begin{array}{l} x_8 = \frac{s_8 - s_f}{s_{fg}} = \frac{6.8000 - 0.6492}{7.4996} = 0.8201 \\ s_8 = s_6 \end{array} \right\}$$

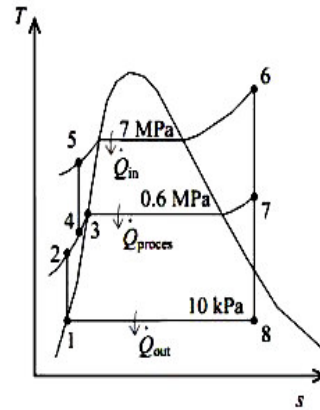
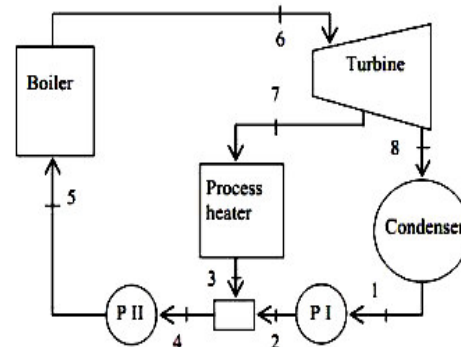
$$h_8 = h_f + x_8 h_{fg} = 191.81 + (0.8201)(2392.1) = 2153.6 \text{ kJ/kg}$$

Then,

$$\dot{W}_{T,out} = \dot{m}_6 (h_6 - h_7) + \dot{m}_8 (h_7 - h_8) \\ = (30 \text{ kg/s})(3411.4 - 2774.6) \text{ kJ/kg} + (22.5 \text{ kg/s})(2774.6 - 2153.6) \text{ kJ/kg} = 33,077 \text{ kW}$$

$$\dot{W}_{p,in} = \dot{m}_1 w_{pl,in} + \dot{m}_4 w_{pll,in} = (22.5 \text{ kg/s})(0.60 \text{ kJ/kg}) + (30 \text{ kg/s})(6.57 \text{ kJ/kg}) = 210.6 \text{ kW}$$

$$\dot{W}_{net} = \dot{W}_{T,out} - \dot{W}_{p,in} = 33,077 - 210.6 = 32,866 \text{ kW}$$





زمان آزمون (دقیقه): تستی: -- تشریحی: ۱۲۰

تعداد سوالات: تستی: ۵ تشریحی: ۵

نام درس: ترمودینامیک ۲

رشته تحصیلی / کد درس: مهندسی مکانیک حرارت و سیالات - مهندسی مکانیک جامدات - مهندسی مکانیک ساخت و تولید - مهندسی مکانیک تبدیل انرژی ۱۳۱۵۰۲۲

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Assumptions 1 Steady operating conditions exist. 2 Helium is an ideal gas with constant specific heats. 3 Kinetic and potential energy changes are negligible.

Properties The properties of helium are $c_p = 5.1926 \text{ kJ/kg}\cdot\text{K}$ and $k = 1.667$ (Table A-2).

Analysis (a) From the isentropic relations,

$$T_{2s} = T_1 \left(\frac{P_2}{P_1} \right)^{(k-1)/k} = (263\text{K})(3)^{0.667/1.667} = 408.2\text{K}$$

$$T_{4s} = T_3 \left(\frac{P_4}{P_3} \right)^{(k-1)/k} = (323\text{K}) \left(\frac{1}{3} \right)^{0.667/1.667} = 208.1\text{K}$$

and

$$\eta_T = \frac{h_3 - h_4}{h_3 - h_{4s}} = \frac{T_3 - T_4}{T_3 - T_{4s}} \longrightarrow T_4 = T_3 - \eta_T (T_3 - T_{4s}) = 323 - (0.80)(323 - 208.1) = 231.1\text{K} = T_{\min}$$

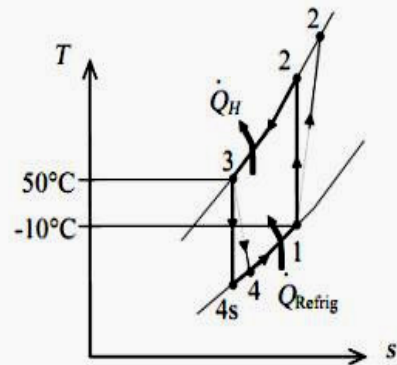
$$\eta_C = \frac{h_{2s} - h_1}{h_2 - h_1} = \frac{T_{2s} - T_1}{T_2 - T_1} \longrightarrow T_2 = T_1 + (T_{2s} - T_1) / \eta_C = 263 + (408.2 - 263) / (0.80) = 444.5\text{K}$$

(b) The COP of this gas refrigeration cycle is determined from

$$\begin{aligned} \text{COP}_R &= \frac{q_L}{w_{\text{net},\text{in}}} = \frac{q_L}{w_{\text{comp},\text{in}} - w_{\text{turb},\text{out}}} \\ &= \frac{h_1 - h_4}{(h_2 - h_1) - (h_3 - h_4)} \\ &= \frac{T_1 - T_4}{(T_2 - T_1) - (T_3 - T_4)} \\ &= \frac{263 - 231.1}{(444.5 - 263) - (323 - 231.1)} = 0.356 \end{aligned}$$

(c) The mass flow rate of helium is determined from

$$\dot{m} = \frac{\dot{Q}_{\text{refrig}}}{q_L} = \frac{\dot{Q}_{\text{refrig}}}{h_1 - h_4} = \frac{\dot{Q}_{\text{refrig}}}{c_p (T_1 - T_4)} = \frac{18 \text{ kJ/s}}{(5.1926 \text{ kJ/kg}\cdot\text{K})(263 - 231.1)\text{K}} = 0.109 \text{ kg/s}$$





کُد سری سؤال: یک (۱)

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Analysis (a) A relation for Δu is obtained from the general relation

$$\Delta u = u_2 - u_1 = \int_{T_1}^{T_2} c_v dT + \int_{v_1}^{v_2} \left(T \left(\frac{\partial P}{\partial T} \right)_v - P \right) dv$$

The equation of state for the specified gas can be expressed as

$$P = \frac{RT}{v-a} \longrightarrow \left(\frac{\partial P}{\partial T} \right)_v = \frac{R}{v-a}$$

Thus,

$$T \left(\frac{\partial P}{\partial T} \right)_v - P = \frac{RT}{v-a} - P = P - P = 0$$

Substituting, $\Delta u = \int_{T_1}^{T_2} c_v dT$

(b) A relation for Δh is obtained from the general relation

$$\Delta h = h_2 - h_1 = \int_{T_1}^{T_2} c_p dT + \int_{P_1}^{P_2} \left(v - T \left(\frac{\partial v}{\partial T} \right)_p \right) dP$$

The equation of state for the specified gas can be expressed as

$$v = \frac{RT}{P} + a \longrightarrow \left(\frac{\partial v}{\partial T} \right)_p = \frac{R}{P}$$

Thus,

$$v - T \left(\frac{\partial v}{\partial T} \right)_p = v - T \frac{R}{P} = v - (v - a) = a$$

Substituting,

$$\Delta h = \int_{T_1}^{T_2} c_p dT + \int_{P_1}^{P_2} a dP = \int_{T_1}^{T_2} c_p dT + a(P_2 - P_1)$$

(c) A relation for Δs is obtained from the general relation

$$\Delta s = s_2 - s_1 = \int_{T_1}^{T_2} \frac{c_p}{T} dT - \int_{P_1}^{P_2} \left(\frac{\partial v}{\partial T} \right)_p dP$$

Substituting $(\partial v / \partial T)_p = R/T$,

$$\Delta s = \int_{T_1}^{T_2} \frac{c_p}{T} dT - \int_{P_1}^{P_2} \left(\frac{R}{P} \right)_p dP = \int_{T_1}^{T_2} \frac{c_p}{T} dT - R \ln \frac{P_2}{P_1}$$

For an isothermal process $dT = 0$ and these relations reduce to

$$\Delta u = 0, \quad \Delta h = a(P_2 - P_1), \quad \text{and} \quad \Delta s = -R \ln \frac{P_2}{P_1}$$



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Properties The inlet and the exit states of the air are completely specified, and the total pressure is 1 atm. The properties of the air at various states are determined from the psychrometric chart (Figure A-31) to be

$$h_1 = 23.5 \text{ kJ/kg dry air}$$

$$\omega_1 = 0.0053 \text{ kg H}_2\text{O/kg dry air} (= \omega_2)$$

$$\nu_1 = 0.809 \text{ m}^3/\text{kg dry air}$$

$$h_3 = 42.3 \text{ kJ/kg dry air}$$

$$\omega_3 = 0.0087 \text{ kg H}_2\text{O/kg dry air}$$

Analysis (a) The amount of moisture in the air remains constant it flows through the heating section ($\omega_1 = \omega_2$), but increases in the humidifying section ($\omega_3 > \omega_2$). The mass flow rate of dry air is

$$\dot{m}_a = \frac{\dot{V}_1}{\nu_1} = \frac{35 \text{ m}^3/\text{min}}{0.809 \text{ m}^3/\text{kg}} = 43.3 \text{ kg/min}$$

Noting that $Q = W = 0$, the energy balance on the humidifying section can be expressed as

$$\dot{E}_{\text{in}} - \dot{E}_{\text{out}} = \Delta \dot{E}_{\text{system}} \stackrel{\approx 0 \text{ (steady)}}{=} 0$$

$$\dot{E}_{\text{in}} = \dot{E}_{\text{out}}$$

$$\sum \dot{m}_i h_i = \sum \dot{m}_e h_e \quad \longrightarrow \quad \dot{m}_w h_w + \dot{m}_a h_2 = \dot{m}_a h_3$$

$$(\omega_3 - \omega_2) h_w + h_2 = h_3$$

Solving for h_2 ,

$$h_2 = h_3 - (\omega_3 - \omega_2) h_{g@100^\circ\text{C}} = 42.3 - (0.0087 - 0.0053)(2675.6) = 33.2 \text{ kJ/kg dry air}$$

Thus at the exit of the heating section we have $\omega_2 = 0.0053 \text{ kg H}_2\text{O/kg dry air}$ and $h_2 = 33.2 \text{ kJ/kg dry air}$, which completely fixes the state. Then from the psychrometric chart we read

$$T_2 = 19.5^\circ\text{C}$$

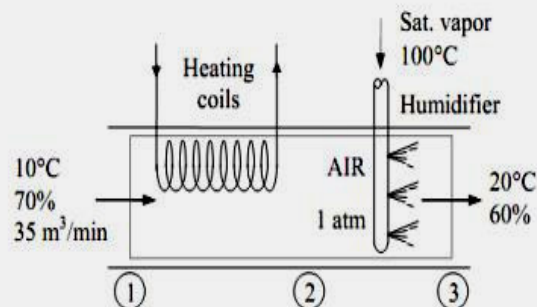
$$\phi_2 = 37.8\%$$

(b) The rate of heat transfer to the air in the heating section is

$$\dot{Q}_{\text{in}} = \dot{m}_a (h_2 - h_1) = (43.3 \text{ kg/min})(33.2 - 23.5) \text{ kJ/kg} = 420 \text{ kJ/min}$$

(c) The amount of water added to the air in the humidifying section is determined from the conservation of mass equation of water in the humidifying section,

$$\dot{m}_w = \dot{m}_a (\omega_3 - \omega_2) = (43.3 \text{ kg/min})(0.0087 - 0.0053) = 0.15 \text{ kg/min}$$





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Assumptions 1 Both the reactants and products are ideal gases. **2** Combustion is complete.

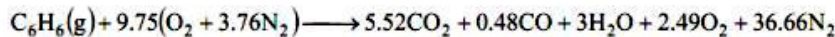
Analysis The theoretical combustion equation of C_6H_6 with stoichiometric amount of air is



where a_{th} is the stoichiometric coefficient and is determined from the O_2 balance,

$$a_{th} = 6 + 1.5 = 7.5$$

Then the actual combustion equation with 30% excess air becomes



The heat transfer for this constant volume combustion process is determined from the energy balance $E_{in} - E_{out} = \Delta E_{system}$ applied on the combustion chamber with $W = 0$. It reduces to

$$-Q_{out} = \sum N_P (\bar{h}_f^\circ + \bar{h} - \bar{h}^\circ - P\bar{v})_P - \sum N_R (\bar{h}_f^\circ + \bar{h} - \bar{h}^\circ - P\bar{v})_R$$

Since both the reactants and the products behave as ideal gases, all the internal energy and enthalpies depend on temperature only, and the $P\bar{v}$ terms in this equation can be replaced by $R_u T$.

It yields

$$-Q_{out} = \sum N_P (\bar{h}_f^\circ + \bar{h}_{1000\text{ K}} - \bar{h}_{298\text{ K}} - R_u T)_P - \sum N_R (\bar{h}_f^\circ - R_u T)_R$$

since the reactants are at the standard reference temperature of 25°C. From the tables,

Substance	\bar{h}_f° kJ/kmol	$\bar{h}_{298\text{ K}}$ kJ/kmol	$\bar{h}_{1000\text{ K}}$ kJ/kmol
$C_6H_6(g)$	82,930	---	---
O_2	0	8682	31,389
N_2	0	8669	30,129
$H_2O(g)$	-241,820	9904	35,882
CO	-110,530	8669	30,355
CO_2	-393,520	9364	42,769

Thus,

$$\begin{aligned} -Q_{out} &= (5.52)(-393,520 + 42,769 - 9364 - 8.314 \times 1000) \\ &\quad + (0.48)(-110,530 + 30,355 - 8669 - 8.314 \times 1000) \\ &\quad + (3)(-241,820 + 35,882 - 9904 - 8.314 \times 1000) \\ &\quad + (2.49)(0 + 31,389 - 8682 - 8.314 \times 1000) \\ &\quad + (36.66)(0 + 30,129 - 8669 - 8.314 \times 1000) \\ &\quad - (1)(82,930 - 8.314 \times 298) - (9.75)(4.76)(-8.314 \times 298) \\ &= -2,200,433 \text{ kJ} \end{aligned}$$

or $Q_{out} = 2,200,433 \text{ kJ}$