

HOMEWORK #4 SOLUTIONS

Problem #1: RDK 18.24

Solve: (a) The mean free path is

$$\lambda = \frac{1}{4\sqrt{2}\pi(N/V)r^2}$$

where $r \approx 0.5 \times 10^{-10}$ m is the atomic radius for helium and N/V is the gas number density. From the ideal-gas law,

$$\begin{aligned} \frac{N}{V} &= \frac{p}{kT} = \frac{0.10 \text{ atm} \times 101,300 \text{ Pa/atm}}{(1.38 \times 10^{-23} \text{ J/K})(10 \text{ K})} = 7.34 \times 10^{25} \text{ m}^{-3} \\ \Rightarrow \lambda &= \frac{1}{4\sqrt{2}\pi(7.34 \times 10^{25} \text{ m}^{-3})(0.5 \times 10^{-10} \text{ m})^2} = 3.1 \times 10^{-7} \text{ m} = 310 \text{ nm} \end{aligned}$$

(b) The root-mean-square speed is

$$v_{\text{rms}} = \sqrt{\frac{3k_{\text{B}}T}{m}} = \sqrt{\frac{3(1.38 \times 10^{-23} \text{ J/K})(10 \text{ K})}{4 \times (1.661 \times 10^{-27} \text{ kg})}} = 250 \text{ m/s}$$

where we used $A = 4$ u as the atomic mass of helium.

(c) The average energy per atom is $e_{\text{avg}} = \frac{3}{2}k_{\text{B}}T = \frac{3}{2}(1.38 \times 10^{-23} \text{ J/K})(10 \text{ K}) = 2.1 \times 10^{-22} \text{ J}$.

Problem #2: RDK 18.32

Solve: The conservation of energy equation $(\Delta E_{\text{th}})_{\text{gas}} + (\Delta E_{\text{th}})_{\text{solid}} = 0$ J is

$$\begin{aligned} n_{\text{gas}}(C_V)_{\text{gas}}(T_f - T)_{\text{gas}} + n_{\text{solid}}(C_V)_{\text{solid}}(T_f - T_i)_{\text{solid}} &= 0 \text{ J} \\ \Rightarrow (1.0 \text{ mol})(12.5 \text{ J/mol K})(-50 \text{ K}) + (1.0 \text{ mol})(25.0 \text{ J/mol K})(\Delta T)_{\text{solid}} &= 0 \Rightarrow (\Delta T)_{\text{solid}} = 25^\circ\text{C} \end{aligned}$$

The temperature of the solid increases by 25°C .

Problem #3: RDK 18.34

Visualize: At low temperatures, $C_V = \frac{3}{2}R = 12.5 \text{ J/mol K}$. At room temperature and modestly hot temperatures,

$C_V = \frac{5}{2}R = 20.8 \text{ J/mol K}$. At very hot temperatures, $C_V = \frac{7}{2}R = 29.1 \text{ J/mol K}$.

Solve: (a) The number of moles of diatomic hydrogen gas in the rigid container is

$$\frac{0.20 \text{ g}}{2 \text{ g/mol}} = 0.10 \text{ mol}$$

The heat needed to change the temperature of the gas from 50 K to 100 K at constant volume is

$$Q = \Delta E_{\text{th}} = nC_V\Delta T = (0.10 \text{ mol})(12.5 \text{ J/mol K})(100 \text{ K} - 50 \text{ K}) = 62 \text{ J}$$

(b) To raise the temperature from 250 K to 300 K,

$$Q = \Delta E_{\text{th}} = (0.10 \text{ mol})(20.8 \text{ J/mol K})(300 \text{ K} - 250 \text{ K}) = 100 \text{ J}$$

(c) To raise the temperature from 2250 K to 2300 K, $Q = \Delta E_{\text{th}} = nC_V\Delta T = (0.10 \text{ mol})(29.1 \text{ J/mol K})(50 \text{ K}) = 150 \text{ J}$

Problem #4: RDK 18.48

Solve: (a) The number of molecules of helium is

$$N_{\text{helium}} = \frac{pV}{k_B T} = \frac{(2.0 \times 1.013 \times 10^5 \text{ Pa})(100 \times 10^{-6} \text{ m}^3)}{(1.38 \times 10^{-23} \text{ J/K})(373 \text{ K})} = 3.936 \times 10^{21}$$

$$\Rightarrow n_{\text{helium}} = \frac{3.936 \times 10^{21}}{6.022 \times 10^{23} \text{ mol}^{-1}} = 6.536 \times 10^{-3} \text{ mol}$$

The initial internal energy of helium is

$$E_{\text{helium } i} = \frac{3}{2} N_{\text{helium}} k_B T = 30.4 \text{ J} \approx 30 \text{ J}$$

The number of molecules of argon is

$$N_{\text{argon}} = \frac{pV}{k_B T} = \frac{(4.0 \times 1.013 \times 10^5 \text{ Pa})(200 \times 10^{-6} \text{ m}^3)}{(1.38 \times 10^{-23} \text{ J/K})(673 \text{ K})} = 8.726 \times 10^{21}$$

$$\Rightarrow n_{\text{argon}} = \frac{8.726 \times 10^{21}}{6.022 \times 10^{23} \text{ mol}^{-1}} = 1.449 \times 10^{-2} \text{ mol}$$

The initial thermal energy of argon is

$$E_{\text{argon } i} = \frac{3}{2} N_{\text{argon}} k_B T = 121.6 \text{ J} \approx 122 \text{ J}$$

(b) The equilibrium condition for monatomic gases is

$$(\epsilon_{\text{helium } f})_{\text{avg}} = (\epsilon_{\text{argon } f})_{\text{avg}} = (\epsilon_{\text{total}})_{\text{avg}}$$

$$\Rightarrow \frac{E_{\text{helium } f}}{n_{\text{helium}}} = \frac{E_{\text{argon } f}}{n_{\text{argon}}} = \frac{E_{\text{tot}}}{n_{\text{tot}}} = \frac{(30.4 + 121.6) \text{ J}}{(6.54 \times 10^{-3} + 1.449 \times 10^{-2}) \text{ mol}} = 7228 \text{ J/mol}$$

$$\Rightarrow E_{\text{helium } f} = (7228 \text{ J/mol})n_{\text{helium}} = (7228 \text{ J/mol})(6.54 \times 10^{-3} \text{ mol}) = 47.3 \text{ J} \approx 47 \text{ J}$$

$$E_{\text{argon } f} = (7228 \text{ J/mol})n_{\text{argon}} = (7228 \text{ J/mol})(1.449 \times 10^{-2} \text{ mol}) = 104.7 \text{ J} \approx 105 \text{ J}$$

(c) The amount of heat transferred is

$$E_{\text{helium } f} - E_{\text{helium } i} = 47.3 \text{ J} - 30.4 \text{ J} = 16.9 \text{ J} \quad E_{\text{argon } f} - E_{\text{argon } i} = 104.7 \text{ J} - 121.6 \text{ J} = -16.9 \text{ J}$$

The helium gains 16.9 J of heat energy and the argon loses 16.9 J. Thus approximately 17 J are transferred from the argon to the helium.

(d) The equilibrium condition for monatomic gases is

$$(\epsilon_{\text{helium}})_{\text{avg}} = (\epsilon_{\text{argon}})_{\text{avg}} \Rightarrow \frac{E_{\text{helium } f}}{N_{\text{helium}}} = \frac{E_{\text{argon } f}}{N_{\text{argon}}} = \frac{3}{2} k_B T_f$$

Substituting the above values,

$$\frac{47.3 \text{ J}}{3.936 \times 10^{21}} = \frac{104.7 \text{ J}}{8.726 \times 10^{21}} = \frac{3}{2} (1.38 \times 10^{-23} \text{ J/K}) T_f \Rightarrow T_f = 580 \text{ K} = 307^\circ \text{C}$$

(e) The final pressure of the helium and argon are

$$p_{\text{helium } f} = \frac{N_{\text{helium}} k_B T}{V_{\text{helium}}} = \frac{(3.936 \times 10^{21})(1.38 \times 10^{-23} \text{ J/K})(580 \text{ K})}{100 \times 10^{-6} \text{ m}^3} = 3.15 \times 10^5 \text{ Pa} \approx 3.1 \text{ atm}$$

$$p_{\text{argon } f} = \frac{N_{\text{argon}} k_B T}{V_{\text{argon}}} = \frac{(8.726 \times 10^{21})(1.38 \times 10^{-23} \text{ J/K})(580 \text{ K})}{200 \times 10^{-6} \text{ m}^3} = 3.49 \times 10^5 \text{ Pa} \approx 3.4 \text{ atm}$$

Problem #5: RDK 19.2

Solve: During each cycle, the work done by the engine is $W_{\text{out}} = 200 \text{ J}$ and the engine exhausts $Q_{\text{C}} = 400 \text{ J}$ of heat energy. By conservation of energy,

$$Q_{\text{H}} = W_{\text{out}} + Q_{\text{C}} = 200 \text{ J} + 400 \text{ J} = 600 \text{ J}$$

Thus, the efficiency of the engine is

$$\eta = \frac{W_{\text{out}}}{Q_{\text{H}}} = \frac{200 \text{ J}}{600 \text{ J}} = 0.33$$

Problem #6: RDK 19.14

Model: The heat engine follows a closed cycle, starting and ending in the original state. The cycle consists of three individual processes.

Solve: (a) The work done by the heat engine per cycle is the area enclosed by the p -versus- V graph. We get

$$W_{\text{out}} = \frac{1}{2}(200 \text{ kPa})(100 \times 10^{-6} \text{ m}^3) = 10 \text{ J}$$

The heat energy transferred into the engine is $Q_{\text{H}} = 30 \text{ J} + 84 \text{ J} = 114 \text{ J}$. Because $W_{\text{out}} = Q_{\text{H}} - Q_{\text{C}}$, the heat energy exhausted is

$$Q_{\text{C}} = Q_{\text{H}} - W_{\text{out}} = 114 \text{ J} - 10 \text{ J} = 104 \text{ J} \approx 0.10 \text{ kJ}$$

(b) The thermal efficiency of the engine is

$$\eta = \frac{W_{\text{out}}}{Q_{\text{H}}} = \frac{10 \text{ J}}{114 \text{ J}} = 0.088$$

Assess: Practical engines have thermal efficiencies in the range $\eta \approx 0.1 - 0.4$.

Problem #7: RDK 19.17

Model: The Brayton cycle involves two adiabatic processes and two isobaric processes. The adiabatic processes involve compression and expansion through the turbine.

Solve: The thermal efficiency for the Brayton cycle is $\eta_{\text{B}} = 1 - r_{\text{p}}^{(1-\gamma)/\gamma}$, where $\gamma = C_{\text{p}}/C_{\text{v}}$ and r_{p} is the pressure ratio. For a diatomic gas $\gamma = 1.4$. For an adiabatic process,

$$p_1 V_1^\gamma = p_2 V_2^\gamma \Rightarrow p_2/p_1 = (V_1/V_2)^\gamma$$

Because the volume is halved, $V_2 = \frac{1}{2}V_1$ so

$$r_{\text{p}} = p_2/p_1 = (2)^\gamma = 2^{1.4} = 2.639$$

The efficiency is

$$\eta_{\text{B}} = 1 - (2.639)^{-0.4/1.4} = 0.24$$

Problem #8: RDK 19.53

Model: The heat engine follows a closed cycle. For a diatomic gas, $C_V = \frac{5}{2}R$ and $C_P = \frac{7}{2}R$.

Visualize: Please refer to Figure P19.53.

Solve: (a) Since $T_1 = 293$ K, the number of moles of the gas is

$$n = \frac{p_1 V_1}{RT_1} = \frac{(0.5 \times 1.013 \times 10^5 \text{ Pa})(10 \times 10^{-6} \text{ m}^3)}{(8.31 \text{ J/mol K})(293 \text{ K})} = 2.08 \times 10^{-4} \text{ mol}$$

At point 2, $V_2 = 4V_1$ and $p_2 = 3p_1$. The temperature is calculated as follows:

$$\frac{p_1 V_1}{T_1} = \frac{p_2 V_2}{T_2} \Rightarrow T_2 = \frac{p_2 V_2}{p_1 V_1} T_1 = (3)(4)(293 \text{ K}) = 3516 \text{ K}$$

At point 3, $V_3 = V_2 = 4V_1$ and $p_3 = p_1$. The temperature is calculated as before:

$$T_3 = \frac{p_3 V_3}{p_1 V_1} T_1 = (1)(4)(293 \text{ K}) = 1172 \text{ K}$$

For process $1 \rightarrow 2$, the work done is the area under the p -versus- V curve. That is,

$$\begin{aligned} W_s &= (0.5 \text{ atm})(40 \text{ cm}^3 - 10 \text{ cm}^3) + \frac{1}{2}(1.5 \text{ atm} - 0.5 \text{ atm})(40 \text{ cm}^3 - 10 \text{ cm}^3) \\ &= (30 \times 10^{-6} \text{ m}^3)(1 \text{ atm}) \left(\frac{1.013 \times 10^5 \text{ Pa}}{1 \text{ atm}} \right) = 3.04 \text{ J} \end{aligned}$$

The change in the thermal energy is

$$\Delta E_{\text{th}} = nC_V \Delta T = (2.08 \times 10^{-4} \text{ mol}) \frac{5}{2} (8.31 \text{ J/mol K})(3516 \text{ K} - 293 \text{ K}) = 13.93 \text{ J}$$

The heat is $Q = W_s + \Delta E_{\text{th}} = 16.97$ J. For process $2 \rightarrow 3$, the work done is $W_s = 0$ J and

$$\begin{aligned} Q &= \Delta E_{\text{th}} = nC_V \Delta T = n \left(\frac{5}{2} R \right) (T_3 - T_2) \\ &= (2.08 \times 10^{-4} \text{ mol}) \frac{5}{2} (8.31 \text{ J/mol K})(1172 \text{ K} - 3516 \text{ K}) = -10.13 \text{ J} \end{aligned}$$

For process $3 \rightarrow 1$,

$$\begin{aligned} W_s &= (0.5 \text{ atm})(10 \text{ cm}^3 - 40 \text{ cm}^3) = (0.5 \times 1.013 \times 10^5 \text{ Pa})(-30 \times 10^{-6} \text{ m}^3) = -1.52 \text{ J} \\ \Delta E_{\text{th}} &= nC_V \Delta T = (2.08 \times 10^{-4} \text{ mol}) \frac{5}{2} (8.31 \text{ J/mol K})(293 \text{ K} - 1172 \text{ K}) = -3.80 \text{ J} \end{aligned}$$

The heat is $Q = \Delta E_{\text{th}} + W_s = -5.32$ J.

	W_s (J)	Q (J)	ΔE_{th}
$1 \rightarrow 2$	3.04	16.97	13.93
$2 \rightarrow 3$	0	-10.13	-10.13
$3 \rightarrow 1$	-1.52	-5.32	-3.80
Net	1.52	1.52	0

(b) The efficiency of the engine is

$$\eta = \frac{W_{\text{net}}}{Q_{\text{H}}} = \frac{1.52 \text{ J}}{16.97 \text{ J}} = 0.090 = 9.0\%$$

(c) The power output of the engine is

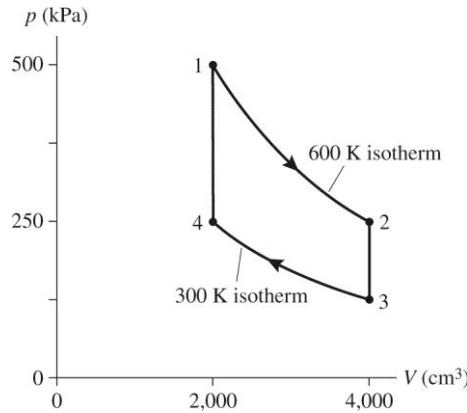
$$500 \left(\frac{\text{revolutions}}{\text{min}} \right) \left(\frac{1 \text{ min}}{60 \text{ s}} \right) \left(\frac{W_{\text{net}}}{\text{revolution}} \right) = \left(\frac{500}{60} \right) (1.52 \text{ J/s}) = 13 \text{ W}$$

Assess: Note that more than two significant figures are retained in part (a) because the results are intermediate. For a closed cycle, as expected, $(W_s)_{\text{net}} = Q_{\text{net}}$ and $(\Delta E_{\text{th}})_{\text{net}} = 0 \text{ J}$.

Problem #9: RDK 19.63 (BONUS)

Model: The closed cycle of the heat engine involves the following four processes: isothermal expansion, isochoric cooling, isothermal compression, and isochoric heating. For a monatomic gas $C_V = \frac{3}{2}R$.

Visualize:



Solve: Using the ideal-gas law,

$$p_1 = \frac{nRT_1}{V_1} = \frac{(0.20 \text{ mol})(8.31 \text{ J/mol K})(600 \text{ K})}{2.0 \times 10^{-3} \text{ m}^3} = 4.986 \times 10^5 \text{ Pa}$$

At point 2, because of the isothermal conditions, $T_2 = T_1 = 600 \text{ K}$ and

$$p_2 = p_1 \frac{V_1}{V_2} = (4.986 \times 10^5 \text{ Pa}) \left(\frac{2.0 \times 10^{-3} \text{ m}^3}{4.0 \times 10^{-3} \text{ m}^3} \right) = 2.493 \times 10^5 \text{ Pa}$$

At point 3, because it is an isochoric process, $V_3 = V_2 = 4000 \text{ cm}^3$ and

$$p_3 = p_2 \frac{T_3}{T_2} = (2.493 \times 10^5 \text{ Pa}) \left(\frac{300 \text{ K}}{600 \text{ K}} \right) = 1.247 \times 10^5 \text{ Pa}$$

Likewise at point 4, $T_4 = T_3 = 300 \text{ K}$ and

$$p_4 = p_3 \frac{V_3}{V_4} = (1.247 \times 10^5 \text{ Pa}) \left(\frac{4.0 \times 10^{-3} \text{ m}^3}{2.0 \times 10^{-3} \text{ m}^3} \right) = 2.493 \times 10^5 \text{ Pa}$$

Let us now calculate $W_{\text{net}} = W_{1 \rightarrow 2} + W_{2 \rightarrow 3} + W_{3 \rightarrow 4} + W_{4 \rightarrow 1}$. For the isothermal processes,

$$W_{1 \rightarrow 2} = nRT_1 \ln \frac{V_2}{V_1} = (0.20 \text{ mol})(8.31 \text{ J/mol K})(600 \text{ K}) \ln(2) = 691.2 \text{ J}$$

$$W_{3 \rightarrow 4} = nRT_3 \ln \frac{V_4}{V_3} = (0.20 \text{ mol})(8.31 \text{ J/mol K})(300 \text{ K}) \ln\left(\frac{1}{2}\right) = -345.6 \text{ J}$$

For the isochoric processes, $W_{2 \rightarrow 3} = W_{4 \rightarrow 1} = 0 \text{ J}$. Thus, the work done per cycle is $W_{\text{net}} = 345.6 \text{ J} \approx 350 \text{ J}$. Because $Q = W_s + \Delta E_{\text{th}}$,

$$Q_{1 \rightarrow 2} = W_{1 \rightarrow 2} + (\Delta E_{\text{th}})_{1 \rightarrow 2} = 691.2 \text{ J} + 0 \text{ J} = 691.2 \text{ J}$$

For the first isochoric process,

$$\begin{aligned} Q_{2 \rightarrow 3} &= nC_V \Delta T = (0.20 \text{ mol}) \left(\frac{3}{2} R \right) (T_3 - T_2) \\ &= (0.20 \text{ mol}) \frac{3}{2} (8.31 \text{ J/mol K}) (300 \text{ K} - 600 \text{ K}) = -747.9 \text{ J} \end{aligned}$$

For the second isothermal process

$$Q_{3 \rightarrow 4} = W_{3 \rightarrow 4} + (\Delta E_{\text{th}})_{3 \rightarrow 4} = -345.6 \text{ J} + 0 \text{ J} = -345.6 \text{ J}$$

For the second isochoric process,

$$\begin{aligned} Q_{4 \rightarrow 1} &= nC_V \Delta T = n \left(\frac{3}{2} R \right) (T_1 - T_4) \\ &= (0.20 \text{ mol}) \left(\frac{3}{2} \right) (8.31 \text{ J/mol K}) (600 \text{ K} - 300 \text{ K}) = 747.9 \text{ J} \end{aligned}$$

Thus, $Q_H = Q_{1 \rightarrow 2} + Q_{4 \rightarrow 1} = 1439.1 \text{ J}$. The thermal efficiency of the engine is

$$\eta = \frac{W_{\text{net}}}{Q_H} = \frac{345.6 \text{ J}}{1439.1 \text{ J}} = 0.24 = 24\%$$