

①

Ch. 19

Q6. (a) ΔE_{int} depends only on initial & final state.
 $\rightarrow$ path independent.

$$\therefore 1 = 2 = 3 = 4$$

(b) $W = \int_{V_i}^{V_f} p dV \rightarrow$ area below path.

$$\therefore 4 > 3 > 2 > 1$$

(c) $\Delta E_{\text{int}} = Q - W$

$$\rightarrow Q = \Delta E_{\text{int}} + W$$

Since ΔE_{int} is the same for all paths,

$$4 > 3 > 2 > 1$$

71.E.

Along path A,

$$W_A = \int_1^4 p dV = 40 \int_1^4 dV = 120 \text{ J}$$

Along path B,

$$W_B = \int_1^4 p dV = \int_1^4 (50 - 10V) dV = 50 \cdot (4 - 1) - \frac{10}{2} (4^2 - 1^2) = 75 \text{ J}$$

($\because$ Path B is a straight line passing through two points $(1 \text{ m}^3, 40 \text{ Pa})$ & $(4 \text{ m}^3, 10 \text{ Pa})$.)

We can write $p = a + bV$ and find a & b .)

Along path C,

$$W_C = \int_1^4 p dV = 10 \int_1^4 dV = 30 \text{ J}$$

(2)

76E. Since ΔE_{int} depends only on states (initial & final), in ~~the~~ cyclical processes $\Delta E_{\text{int}} = 0$. (if we consider a complete cycle.)

$$\therefore 0 = Q - W$$

$$\rightarrow Q = W$$

Along $A \rightarrow B$,

$$W_{AB} = \int_1^4 p dV = \int_1^4 \left(\frac{10}{3} + \frac{20}{3} V \right) dV$$

$$= 60 \text{ J}$$

Along $B \rightarrow C$

$$W_{BC} = \int_4^1 p dV = 30 \int_4^1 dV = -90 \text{ J}$$

$$\text{Along } C \rightarrow A, \quad W_{CA} = \int_1^1 p dV = 0$$

$$\therefore Q = W = 60 \text{ J} - 90 \text{ J} + 0 \text{ J} = -30 \text{ J}$$

$\rightarrow$ the gas loses 30 J of energy in the form of heat.

Ch. 20

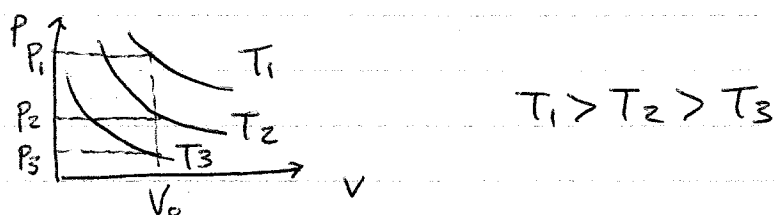
Q1. The pressure of the gas increased but less than doubled.

$$P_1 V = nR(293\text{K})$$

$$P_2 V = nR(313\text{K})$$

$$\rightarrow \frac{P_2}{P_1} = \frac{313}{293} < 2.$$

Q6.



$$\frac{PV}{T} = nR = \text{constant}.$$

At any given volume V_0 , we have

$$\frac{P_1 V_0}{T_1} = \frac{P_2 V_0}{T_2} = \frac{P_3 V_0}{T_3} = \text{constant}.$$

If $T_1 > T_2 > T_3$, then $P_1 > P_2 > P_3$ as we expect from the above graph.

$\therefore$ points 1, 2, 3, 4 are in lower temperature.

Q11.

(a) Same

(b) $\frac{PV}{T} = \text{constant}.$

increases

(c) For adiabatic processes, $PV^\gamma = \text{constant}.$
where $\gamma = \frac{C_p}{C_v} > 1.$

$\therefore P$ decreases as V increases.

(d) $\frac{PV}{T} = \text{const.} \Rightarrow T$ increases.

(4)

Q13. ΔE_{int} along path abc and along ac are the same, since initial state & final state are the same for each paths.

Path ab is an isothermal process.

Since $\Delta E_{int} = nC_v \Delta T$, $\Delta E_{int} = 0$ for path ab.

Path ac is adiabatic. So $Q = 0$.

Then $\Delta E_{int} = -W$ ← first law of thermodynamics.

$\therefore \Delta E_{int} = -4J$.

⇒ Overall $\Delta E_{int} = -4J$

Q15. (a) For adiabatic processes,
 $pV^\gamma = \text{const.}$ where $\gamma = \frac{C_p}{C_v}$

$$\gamma_{\text{mono}} = \frac{5R/2}{3R/2} = 1.67$$

$$\gamma_{\text{dia}} = \frac{7R/2}{5R/2} = 1.4$$

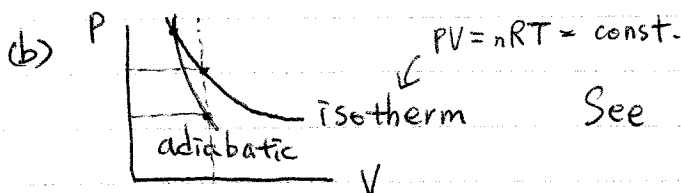
$$\gamma_{\text{poly}} = \frac{4R}{3R} = 1.3$$

⇒ $\gamma_{\text{mono}} > \gamma_{\text{dia}} > \gamma_{\text{poly}}$.

which means for a given volume change,

$$P_{\text{mono}} < P_{\text{dia}} < P_{\text{poly}}$$

Therefore, 1 = polyatomic, 2 = diatomic, 3 = monoatomic.



See Fig. 20-14

P_{isotherm} would be more than $P_{\text{adiabatic}}$. { adiabatic graph is more steeper because $PV^\gamma = \text{const.}$ & $\gamma > 1$.

(5)

5P. Since the molar mass of the ink is $M = 18 \text{ g/mol}$, the number of molecules in $m = 1 \mu\text{g} = 10^{-6} \text{ g}$ of ink is

$$N = \frac{m N_A}{M} = \frac{(10^{-6} \text{ g}) \cdot (6.02 \times 10^{23} \text{ mol}^{-1})}{18 \text{ g} \cdot \text{mol}^{-1}}$$

$$\approx 3 \times 10^{16}$$

The number of creatures in our galaxy, with the assumption made in the problem, is about

$$N' = (5 \times 10^9) \cdot (10^{11}) = 5 \times 10^{20}$$

So the statement was wrong by a factor of about 20,000

11E.

$$PV = nRT$$

$$P_1 V_1 = nRT_1$$

$$P_2 V_2 = nRT_2$$

$$\rightarrow \frac{P_2 V_2}{P_1 V_1} = \frac{T_2}{T_1} \quad \therefore P_2 = \frac{V_1}{V_2} \cdot \frac{T_2}{T_1} \cdot P_1$$

$$P_2 = \frac{1000 \text{ in}^3}{1020 \text{ in}^3} \cdot \frac{300 \text{ K}}{273 \text{ K}} \cdot 24 \text{ lb/in}^2$$

$$= 27.0 \text{ lb/in}^2$$

⑥

18P. (a) $PV = nRT$

$$n = \frac{(2.5 \text{ kPa}) \cdot (1.0 \text{ m}^3)}{(8.31 \text{ J/mol} \cdot \text{K}) \cdot (200 \text{ K})} = 1.5 \text{ mol}$$

(b) $T_B = \frac{(7.5 \text{ kPa}) \cdot (3.0 \text{ m}^3)}{(1.5 \text{ mol}) \cdot (8.31 \text{ J/mol} \cdot \text{K})}$

(c) $T_C = \frac{(2.5 \text{ kPa}) \cdot (3.0 \text{ m}^3)}{(1.5 \text{ mol}) \cdot (8.31 \text{ J/mol} \cdot \text{K})} = 6.0 \times 10^2 \text{ K}$

(d) $\Delta E_{\text{int}} = Q - W$

$\Delta E_{\text{int}} = 0$ for the cyclic process abca.

$Q = W = \frac{1}{2} (V_c - V_a) (P_b - P_c) \leftarrow \text{area of triangle abc}$
 $= \frac{1}{2} \cdot 2 \cdot 5 \times 10^3 = 5 \times 10^3 \text{ (J)}$

($\therefore W = \underbrace{\int_{V_a}^{V_b} p dV}_{\oplus} + 0 + \underbrace{\int_{V_c}^{V_a} p dV}_{\ominus}$
 $= \underbrace{\int_{V_a}^{V_b} p dV}_{\oplus} - \underbrace{\int_{V_a}^{V_c} p dV}_{\oplus} = \text{area of triangle abc.})$

31E. $v_{\text{rms}} = \sqrt{\frac{3RT}{M}}$, M : molar mass

$$\sqrt{\frac{3RT_{\text{H}_2}}{M_{\text{H}_2}}} = \sqrt{\frac{3RT_{\text{He}}}{M_{\text{He}}}}$$

$\Rightarrow T_{\text{He}} = \frac{M_{\text{He}}}{M_{\text{H}_2}} \cdot T_{\text{H}_2} = \frac{4 \text{ g/mol}}{2 \text{ g/mol}} \cdot (20 + 273)$

$= 586 \text{ K}$

$= 313^\circ \text{C}$

(7)

37E.

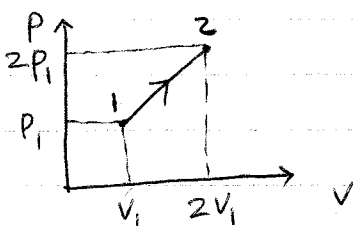
$$U = mgh$$

$$\overline{K} = \frac{3}{2} kT$$

$$\Rightarrow \frac{U}{\overline{K}} = \frac{2mgh}{3kT} = \frac{2 \cdot (0.032 \text{ kg/NA}) \cdot (9.8 \text{ m/s}^2)(0.1 \text{ m})}{3 \cdot (R/\text{NA}) \cdot (273 \text{ K})}$$

$$= 9.2 \times 10^{-6} \quad \leftarrow k = \frac{R}{\text{NA}}$$

64P.



And it is given that

$$P_1 V_1 = nRT_1 \quad (*)$$

(a) The straight line can be expressed as

$$P - P_1 = \frac{2P_1 - P_1}{2V_1 - V_1} (V - V_1)$$

$$\Rightarrow P = \frac{P_1}{V_1} V$$

Using (*), we have

$$P = \frac{nRT_1}{V_1^2} \cdot V$$

$$\text{Now, } W = \int_{V_1}^{2V_1} P dV = \frac{nRT_1}{V_1^2} \int_{V_1}^{2V_1} V dV = \frac{3}{2} nRT_1$$

(b) At point 2, $(2P_1) \cdot (2V_1) = nRT_2$

$$\Delta E_{\text{int}} = \frac{3}{2} nRT_2 - \frac{3}{2} nRT_1$$

$$= \frac{3}{2} \cdot 4P_1 V_1 - \frac{3}{2} nRT_1 = \frac{3}{2} \cdot 4nRT_1 - \frac{3}{2} nRT_1$$

$$= \frac{9}{2} nRT_1$$

(c) $Q = \Delta E_{\text{int}} + W$

$$= \frac{9}{2} nRT_1 + \frac{3}{2} nRT_1 = 6nRT_1$$

(8)

(d) The molar specific heat would be

$$\begin{aligned}
 C &= \frac{Q}{n\Delta T} = \frac{6nRT_1}{n(T_2 - T_1)} \\
 &= \frac{6RT_1}{\frac{(2P_1)(2V_1)}{nR} - T_1} \\
 &= \frac{6RT_1}{4T_1 - T_1} = 2R
 \end{aligned}$$

70 P. (a) For constant pressure process,

$$Q = nC_p\Delta T$$

For a diatomic ideal gas, $C_p = \frac{7}{2}R$.

$$\begin{aligned}
 \therefore Q &= n\frac{7}{2}R\Delta T = \frac{7}{2} \cdot (4\text{ mol}) \cdot (8.31\text{ J/mol}\cdot\text{K}) \cdot (60\text{ K}) \\
 &= 6.98 \times 10^3\text{ J}
 \end{aligned}$$

(b) For any process, $\Delta E_{\text{int}} = nC_v\Delta T$

For a diatomic gas, $C_v = \frac{5}{2}R$

$$\begin{aligned}
 \therefore \Delta E_{\text{int}} &= \frac{5}{2} \cdot (4\text{ mol}) \cdot (8.31\text{ J/mol}\cdot\text{K}) \cdot (60\text{ K}) \\
 &= 4.99 \times 10^3\text{ J}
 \end{aligned}$$

$$\begin{aligned}
 \text{(c)} \quad W &= Q - \Delta E_{\text{int}} \quad (\text{first law of thermodynamics}) \\
 &= 1.99 \times 10^3\text{ J}
 \end{aligned}$$

$$\text{(d)} \quad \Delta K = \frac{3}{2}nR\Delta T = \frac{3}{2} \cdot (4\text{ mol}) \cdot (8.31\text{ J/mol}\cdot\text{K}) \cdot (60\text{ K}) = 299 \times 10^3\text{ J}$$

even if we are dealing with a diatomic gas, ^{since} we want to find "translational" kinetic energy, the factor is $3/2$.

(9)

82P. (a) Isothermal. ($T_1 = T_2$)

$$P_1 V_1 = P_2 V_2 = n R T_1$$

$$P_2 = \frac{V_1}{V_2} P_1 = \frac{1}{4} (32 \text{ atm}) = 8 \text{ atm}$$

$$\begin{aligned} W &= \int_{V_1}^{V_2} p dV = \int_{V_1}^{V_2} \frac{n R T_1}{V} dV \quad \leftarrow \text{constant here} \\ &= P_1 V_1 \ln \frac{V_2}{V_1} = (32 \text{ atm}) \cdot (1.01 \times 10^5 \text{ Pa/atm}) \cdot (1 \times 10^{-3} \text{ m}^3) \ln \frac{4}{1} \\ &= 4.4 \times 10^3 \text{ J} \end{aligned}$$

(b) Adiabatic & Monatomic. ($Q=0$ & $C_V = \frac{3}{2} R$)

$$P_1 V_1^\gamma = P_2 V_2^\gamma$$

$$\Rightarrow P_2 = \left(\frac{V_1}{V_2} \right)^\gamma P_1 = \left(\frac{1}{4} \right)^{5/3} (32 \text{ atm}) = 3.2 \text{ atm}$$

$$\frac{PV}{T} = \text{constant.}$$

$$\Rightarrow T_2 = P_2 V_2 \cdot \frac{T_1}{P_1 V_1} = \frac{(3.2 \text{ atm})(4 \text{ L})}{(32 \text{ atm})(1 \text{ L})} \cdot 300 \text{ K} = 120 \text{ K}$$

$$Q=0 \Rightarrow W = -\Delta E_{\text{int}} = -\frac{3}{2} n R \Delta T$$

$$= -\frac{3}{2} (P_2 V_2 - P_1 V_1)$$

$$= -\frac{3}{2} [(3.2 \text{ atm})(4 \text{ L}) - (32 \text{ atm})(1 \text{ L})] \cdot (1.01 \times 10^5 \text{ Pa/atm}) \cdot (10^{-3} \text{ m}^3/\text{L})$$

$$= 2.9 \times 10^3 \text{ J}$$

(10)

(c) Adiabatic & Diatomic

every procedure in (b) ~~can~~ can be used here. with $\gamma = 1.4$ & $C_v = \frac{5}{2}R$.

$$P_2 = \left(\frac{V_1}{V_2}\right)^\gamma \cdot P_1 = \left(\frac{1L}{4L}\right)^{1.4} \cdot 32 \text{ atm} = 4.6 \text{ atm}$$

$$T_2 = \frac{P_2 V_2}{P_1 V_1} \cdot T_1 = \frac{(4.6 \text{ atm})(4L)}{(32 \text{ atm})(1L)} = 170 \text{ K}$$

$$W = \cancel{Q} - \Delta E_{\text{int}} = -\frac{5}{2} n R \Delta T$$

$$= -\frac{5}{2} (P_2 V_2 - P_1 V_1)$$

$$= 3.4 \times 10^3 \text{ J}$$