

1)

8.314 J/mole · K

$$1 \text{ Pa} = \frac{\text{J}}{\text{m}^3} \quad \text{J} = \text{m}^3 \cdot \text{Pa} \quad 1 \text{ Pa} = 10^{-6} \text{ MPa}$$

$$= \frac{8.314 \text{ Pa} \cdot \text{m}^3}{\text{mole} \cdot \text{K}} = \frac{8.314 \text{ MPa} \cdot \text{cm}^3}{\text{mole} \cdot \text{K}}$$

$$1 \text{ cm}^3 = 1 \text{ mL}$$

$$1 \text{ mL} = 2.6 \times 10^{-4} \text{ gallons}$$

$$1.01 \text{ bar} = 1 \text{ MPa} \quad 14.50 \text{ bar} = 14.5 \text{ MPa}$$

$$145.0 \text{ MPa} = 1 \text{ PSI}$$

$$= \frac{8.314 \text{ cm}^3 \cdot \text{MPa}}{\text{mole} \cdot \text{K}} \left(\frac{1 \text{ mL}}{\text{cm}^3} \right) \left(\frac{2.6 \times 10^{-4} \text{ gal}}{\text{mL}} \right) \left(\frac{145.0 \text{ MPa}}{\text{PSI}} \right) = \frac{1.5 \times 10^{-5} \text{ gal} \cdot \text{PSI}}{\text{mole} \cdot \text{K}}$$

$$\begin{aligned}
 P &= 0.0314 \frac{\text{L} \cdot \text{bar}}{\text{K} \cdot \text{mol}} \\
 T &= 300\text{K} \\
 a &= 1090 \frac{\text{L}^2 \cdot \text{bar} \cdot \text{K}}{\text{mol}^2} \\
 V_1 &= \frac{22.4 \text{ L}}{\text{mol}} \\
 V_2 &= \frac{22.4 \text{ L}}{100}
 \end{aligned}$$

$$\begin{aligned}
 &= -4060 \frac{\text{L} \cdot \text{bar}}{\text{mol}} \\
 &= -4060 \frac{\text{J}}{\text{mol}}
 \end{aligned}$$

$$= - \left[RT \ln \left(\frac{V_2}{V_1} \right) + \frac{7}{2} RT \ln \left(\frac{V_2}{V_1} \right) \right]$$

$$= \sqrt{2} RT - \frac{7}{2} RT$$

$$W = - \int_{V_1}^{V_2} P dV$$

4

$$\Delta 100 \text{ W} = 100 \frac{\text{J}}{\text{s}} \cdot 60 \text{ sec} \cdot \frac{1 \text{ min}}{60 \text{ min}} \cdot \frac{1 \text{ hr}}{24 \text{ hr}} \cdot \frac{1 \text{ day}}{24 \text{ hr}} = 0.69 \times 10^6 \frac{\text{J}}{\text{DAY}}$$

$$\text{C. } 1840 \text{ kcal} \cdot \frac{1 \text{ J}}{0.239 \text{ kcal}} \cdot \frac{1000 \text{ cal}}{1 \text{ kcal}} = 7.70 \times 10^6 \text{ J}$$

$$393 \text{ KJ/mol} \cdot 0.03 \text{ mol} = 32.750 \text{ KJ} = 32.750 \text{ J}$$

$$\frac{19 \text{ cal/mol}}{129 \text{ cal/mol}} = 0.83 \text{ mol}$$

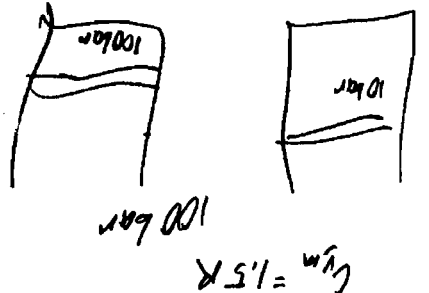
$$\text{B. } (+ O_2 \rightarrow CO_2 \quad 393 \text{ KJ/mol}$$

$$= 490 \text{ KJ} \frac{\text{m}^2}{\text{s}^2} = 490 \text{ J}$$

$$= (50 \text{ kg}) (9.8 \frac{\text{m}}{\text{s}^2}) (1 \text{ m})$$

$$\text{Z. } \Delta PE = mgh$$

2 moles



$P_{ext} = 100 \text{ bar}$

$P_{ext} = 10 \text{ bar}$

Process 1

$$W = - \int_{V_f}^{V_i} P_{ext} dV$$

$$\Delta n = Q + W$$

$$W_i = -P_f \left(\frac{V_f}{nRT} - \frac{V_i}{nRT} \right)$$

$$V_f = \frac{nRT}{P_f}$$

$$V_i = \frac{nRT}{P_i}$$

$$PV = nRT$$

$$W_i = -nRT + nRT \left(\frac{P_i}{P_f} \right)$$

$$W_i = -nRT \left(1 - \frac{P_i}{P_f} \right)$$

$$W_i = -8.314 \text{ J} \cdot 2 \text{ moles} \cdot 300 \text{ K} \left(1 - \frac{10 \text{ bar}}{100 \text{ bar}} \right)$$

$P_f = 100 \text{ bar}$

$P_i = 10 \text{ bar}$

$T = 300 \text{ K}$

$R = 8.314 \text{ J/mole}\cdot\text{K}$

$n = 2$ $T = 300 \text{ K}$

$$W_i = 44895.6 \text{ J} \quad Q = -44895.6 \text{ J}$$

$Q = -W$
 $\Delta n = 0$
 $\Delta H = 0$

reversible process $P_f = 10 \text{ bar}$ $P_i = 100 \text{ bar}$ $\Delta n = 0$

$$W_2 = -nRT \left(1 - \frac{P_i}{P_f} \right)$$

$$W_2 = -2 \text{ moles} \cdot 8.314 \text{ J} \cdot 300 \text{ K} \left(1 - \frac{10 \text{ bar}}{100 \text{ bar}} \right)$$

$$Q = 4489.6 \text{ J}$$

$$W_2 = -4489.6 \text{ J}$$

for isothermal process $nR \ln \frac{V_f}{V_i} = Q = 0$

$\Delta U = 0$ because it is an property of the system and the initial and final conditions are the same

$$q_{total} = q_1 + q_2 = 44,895.6 \text{ J} + (-44,899.5 \text{ J}) = 40,406 \text{ J}$$

$$q_{total} = q_1 + q_2 = -44,895.6 \text{ J} + 44,899.5 \text{ J} = -40,406 \text{ J}$$

for reversible expansion

left to get a constant

$$w = - \int_{V_f}^{V_i} p_{ext} dV = - \int_{V_f}^{V_i} \frac{p}{nRT} dV = - \int_{V_f}^{V_i} \frac{1}{nRT} dV = -nRT \ln \left(\frac{V_i}{V_f} \right)$$

$$\frac{V_f}{p_f} = \frac{V_i}{p_i} \Rightarrow \frac{p_i}{p_f} = \frac{V_i}{V_f} = \frac{\left(\frac{nRT}{p_i} \right)}{\left(\frac{nRT}{p_f} \right)}$$

forward

$$w = -nRT \ln \left(\frac{p_f}{p_i} \right) = -2 \text{ mole} \cdot 8.314 \text{ J} \cdot 300 \text{ K} \ln \left(\frac{10 \text{ bar}}{100 \text{ bar}} \right) = +11,486.2 \text{ J}$$

reverse process

$$q = -11,486.2 \text{ J}$$

$$w = -2 \text{ mole} \cdot 8.314 \text{ J} \cdot 300 \text{ K} \ln \left(\frac{10}{100} \right) = -11,486.2 \text{ J}$$

$$w_{total} = w_{forward} + w_{reverse} = 11,486.2 \text{ J} - 11,486.2 \text{ J} = 0$$

$$q_{total} = q_{forward} + q_{reverse} = -11,486.2 \text{ J} + 11,486.2 \text{ J} = 0$$

$$\Delta U_{total} = 0$$

$$C_p = C_v + nR$$

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$$p = \frac{nR(T - T_0)}{V - V_0}$$

$$C_p = C_v + T \left(\frac{nR}{V} \right) \left(\frac{V - V_0}{T - T_0} \right)$$

$$0 + \frac{d}{nR} = \left(\frac{dT}{dT} \right)$$

$$nR + \frac{d}{V} = 1$$

$$\frac{d}{nR} = 1 - nR$$

$$\frac{(1 - nR)}{nR} = \left(\frac{dT}{dT} \right)$$

$$p = \frac{nR(T - T_0)}{V - V_0}$$

$$p(V - V_0) = nR(T - T_0)$$

C_p & C_v for hard sphere gas that obeys the eqn above

$$C_p = C_v + T \left(\frac{dT}{dT} \right) \left(\frac{dT}{dT} \right)$$

(5)

$$q = \Delta H = \Delta H^\ominus \times n = \Delta H^\ominus \times 200$$

$$q = \Delta H^\ominus \times 200$$

for 200 mole NH_3

stoich coefficient for NH_3

$$q = -10500 \text{ kJ}$$

$$\Delta_r H^\ominus(600) = -105 \times 10^3 \text{ (kJ)} \left(\frac{\text{mole}}{10^3 \text{ J}} \right)$$

$$\Delta_r H^\ominus(600) = -92.22 \text{ kJ/mole} + \frac{\text{kJ}}{\text{mole}} \cdot 1000 \text{ J} + \frac{-45.47 \text{ J}}{\text{mole} \cdot \text{K}} (302 \text{ K})$$

$$\Delta_r H^\ominus(600) = \Delta_r H^\ominus(298 \text{ K}) + \int_{298}^{600} \Delta_r C_p^\ominus dT$$

$$\Delta_r C_p^\ominus = -45.47 \text{ J/mole} \cdot \text{K}$$

$$\Delta_r C_p^\ominus = 2 C_{p,m}^\ominus(\text{NH}_3) - (3 C_{p,m}^\ominus(\text{H}_2) + C_{p,m}^\ominus(\text{N}_2))$$

35.06	28.82	29.13
$2 C_{p,m}^\ominus(\text{NH}_3)$	$3 C_{p,m}^\ominus(\text{H}_2)$	$C_{p,m}^\ominus(\text{N}_2)$

$$\Delta_r C_p^\ominus = 70.12 \text{ J/mole} \cdot \text{K} - 86.46 \text{ J/mole} \cdot \text{K} - 29.13 \text{ J/mole} \cdot \text{K}$$

$$\Delta_r H^\ominus = -92.22 \text{ kJ/mole} @ 298 \text{ K}$$

assume C_p is independent of T

heat of formation of an element in its stable form is 0

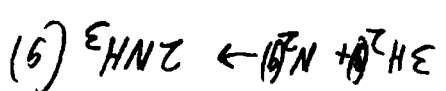
$$\Delta_r H^\ominus = 2(-46.11 \text{ kJ/mole}) - 3(0) + 0$$

$$\Delta_r H^\ominus = \sum \nu \Delta H_f^\ominus(\text{products}) - \sum \nu \Delta H_f^\ominus(\text{reactants})$$

$$\Delta H_f^\ominus \text{H}_2 = 0$$

$$\Delta H_f^\ominus \text{N}_2 = 0$$

$$\Delta H_f^\ominus \text{NH}_3(g) = -46.11 \text{ kJ/mole}$$



7) $\Delta_r H^\ominus$ 298 K 600 K

$$\Delta T = 2.396 \text{ K}$$

$$6550 \text{ J} = 2734 \text{ J/K} \Delta T$$

$$\Delta U = C_v \Delta T$$

Constant Pressure
 $\Delta H \approx \Delta U$
 5/2 value of
 the solid is small

$$\frac{6550 \text{ J} = 2734 \text{ J/K} \Delta T}{\text{Constant Volume}}$$

$$b_u = C_v \Delta T$$

$$\Delta T = 2.396$$

$$C_{p, O_2} = 29.355 \text{ J/K mol} \times 24.9843 = 733.41 \text{ J/K}$$

$$C_{p, CO_2} = 37.11 \text{ J/K mol} \times 0.0167 = 0.62 \text{ J/K}$$

$$+ 2000 \text{ J/K}$$

$$\frac{2734 \text{ J/K}}{2000 \text{ J/K}}$$

- In the container there is 24.9843 mole O_2
 0.0167 mole CO_2

$$0.0167 \times 393 \text{ K} = 6.55 \text{ K}$$

$$12 \text{ g/mole} = \frac{0.0167 \text{ mol Carbon}}{0.29} = 0.0167 \text{ mol Carbon}$$

$$\Delta H_{f, O_2} = 0 \text{ J/mol}$$

$$\Delta H_{f, CO_2} = -393 \text{ J/mol}$$