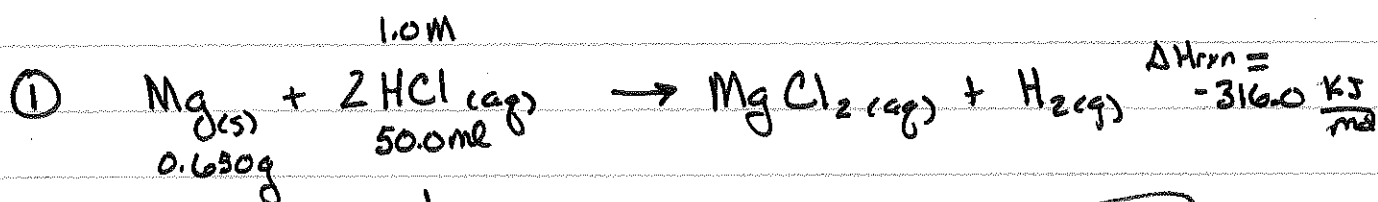


AP Chem Unit 11 Thermodynamics

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Super Problem: Thermodynamics



A) ? Heat released

$$\left(\frac{0.650\text{g Mg}}{1} \right) \left(\frac{1 \text{ mole Mg}}{24.31\text{g Mg}} \right) \left(\frac{316.0 \text{ kJ}}{1 \text{ mole}} \right) = \boxed{8.45 \text{ kJ}}$$

B) $q = mc\Delta T$ $\Delta T = ?$
 $q = 8.45 \text{ kJ} = 8450 \text{ J}$
 $c = 4.18 \text{ J/g}^\circ\text{C}$

$m = \text{mass HCl} + \text{Mg} \Rightarrow 50\text{g} + 0.650\text{g} = 50.7\text{g}$

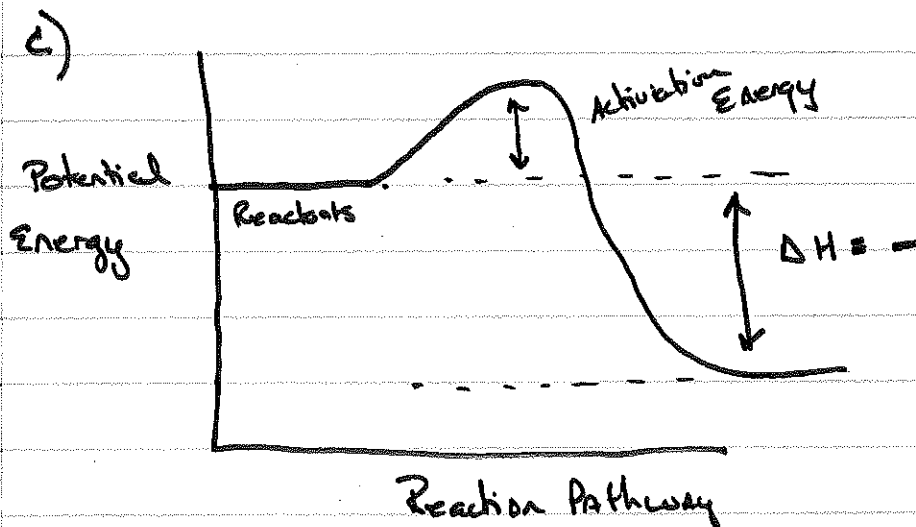
$50\text{ml HCl} \approx 50\text{g}$

$1\text{ml H}_2\text{O} = 1\text{g}$

$$\Delta T = \frac{q}{m c}$$

$$= \frac{8450 \text{ J}}{(50.7\text{g}) \left(4.18 \frac{\text{J}}{\text{g}^\circ\text{C}} \right)}$$

$\Delta T = \boxed{39.90}$





$$V_1 = 650 \text{ mL} \\ 1.0 \text{ atm} \\ \Delta T = 15^\circ\text{C}$$

$$V_2 = 620 \text{ mL}$$

IS TEMP of system ↑ or ↓ ??

The temperature of system is decreasing.

Reason →
must
have

A decrease in volume of gas indicates a decrease in the temperature of the gas the system is at constant pressure and the amount of the gas is not changing

e) Work done? (J)

$$W = -P\Delta V$$

$$P = 1.0 \text{ atm}$$

$$= -(1.0 \text{ atm})(-30 \text{ mL})$$

$$= +30 \text{ atm} \cdot \text{mL}$$

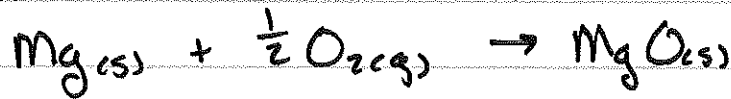
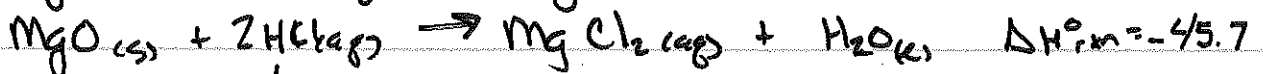
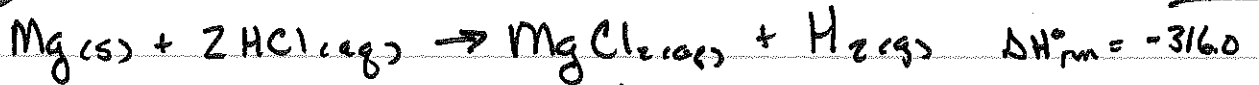
$$\Delta V = V_2 - V_1 = 620 \text{ mL} - 650 \text{ mL} = -30 \text{ mL}$$

want in J

$$\left(\frac{+30 \text{ atm} \cdot \text{mL}}{1} \right) \left(\frac{1 \text{ L}}{1000 \text{ mL}} \right) \left(\frac{101.3 \text{ J}}{1 \text{ L} \cdot \text{atm}} \right) = \boxed{3.04 \text{ J}}$$

f) The statement is wrong. A decrease (-30 mL) in the volume of gas indicates the surroundings are doing work on the system.

NMSI Super Problem: Thermodynamics

g) $\Delta H_{\text{rxn}} = ?$ Given:Soln:

same



R



same



$$\Delta H_{\text{rxn}}^\circ = -556 \frac{\text{kJ}}{\text{mol}}$$



$$\Delta S_{\text{rxn}}^\circ = \sum \Delta S_{\text{p}}^\circ - \sum \Delta S_{\text{r}}^\circ$$

$$= 1 \frac{\text{mol}}{\text{mol rxn}} (27 \frac{\text{J}}{\text{mol}\cdot\text{K}}) - \left[\frac{1 \text{ mol}}{\text{mol rxn}} (33 \frac{\text{J}}{\text{mol}\cdot\text{K}}) + \frac{1 \text{ mole}}{2 \text{ mol rxn}} (205 \frac{\text{J}}{\text{mol}\cdot\text{K}}) \right]$$

$$= 27 \frac{\text{J}}{\text{mol}\cdot\text{K}} - 136 \frac{\text{J}}{\text{mol}\cdot\text{K}}$$

$$\Delta S_{\text{rxn}}^\circ = -109 \frac{\text{J}}{\text{mol}\cdot\text{K}}$$

Super Problem : Thermodynamics

4/5

I) $\Delta G^{\circ}_{\text{Rxn}} = ?$ Mg @ 25° $T = 273 + 25$
 $= 298$

$$\Delta G^{\circ} = \Delta H^{\circ} - T\Delta S^{\circ}$$

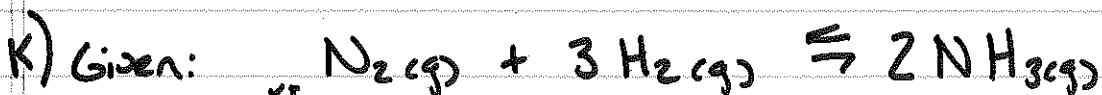
$$= -556 \frac{\text{kJ}}{\text{mol}} - (298 \text{ K}) \left(\frac{-109 \text{ J}}{\text{mol K}} \right) \left(\frac{1 \text{ kJ}}{1000 \text{ J}} \right)$$

$$= -556 \frac{\text{kJ}}{\text{mol}} + 32.5 \frac{\text{kJ}}{\text{mol}}$$

$\Delta G^{\circ} = -524 \text{ kJ/mol}$

J) IS Reaction thermodynamically favored @ 25°C ?

* Since ΔG° is Negative the Reaction is thermodynamically favored



$$\Delta G^{\circ}_{\text{Rxn}} = -34.1 \frac{\text{kJ}}{\text{mol}}$$

$$\Delta H^{\circ}_{\text{Rxn}} = -92.2 \frac{\text{kJ}}{\text{mol}}$$

find: $K = ?$

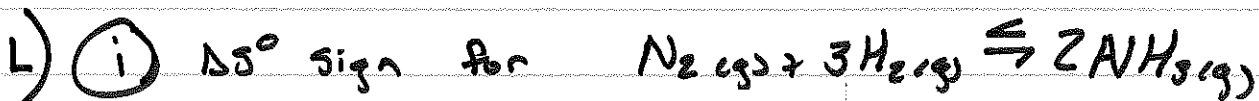
Soln: Since the Reaction is Thermodynamically favored

(i) (ΔG° is Neg) the equilibrium constant will be > 1 as the products are favored

(ii) $\Delta G^{\circ} = -RT \ln K_{\text{eq}} = \frac{(-34.1 \frac{\text{kJ}}{\text{mol}})}{(8.3145 \frac{\text{J}}{\text{mol K}}) (\frac{1 \text{ kJ}}{1000 \text{ J}}) (298 \text{ K})}$

$$\ln K_{\text{eq}} = \frac{-\Delta G^{\circ}}{RT} = 13.8$$

$K_{\text{eq}} = 9.85 \times 10^5$



Soln:

4 moles gas converted to 2 moles gas
 \therefore The Reaction will become less disordered
 (more organized)

$$\therefore -\Delta S$$

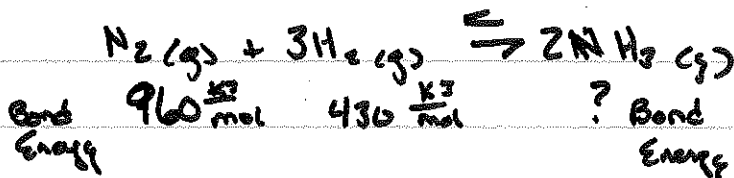
(ii) $\Delta S = ?$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S$$

$$\Delta S = \frac{\Delta H^\circ - \Delta G^\circ}{T} = \frac{-92.2 \frac{\text{kJ}}{\text{mol}} - (-34.1 \frac{\text{kJ}}{\text{mol}})}{298 \text{ K}}$$

$$\Delta S = -195 \frac{\text{kJ}}{\text{mol K}} \quad \text{or} \quad -195 \frac{\text{J}}{\text{mol K}}$$

m) $\Delta H^\circ_{\text{rxn}} = -92.2 \frac{\text{kJ}}{\text{mol}}$



$$\Delta H^\circ_{\text{rxn}} = \sum \text{Bonds Broken} - \sum \text{Bonds Formed} \quad \leftarrow \begin{matrix} \text{coeff} \\ \text{NH}_3 \end{matrix}$$

$$-92.2 \frac{\text{kJ}}{\text{mol}} = [960 \frac{\text{kJ}}{\text{mol}} + 3(430 \frac{\text{kJ}}{\text{mol}})] - [2(3X)]$$

$$-92.2 \frac{\text{kJ}}{\text{mol}} = 2250 \frac{\text{kJ}}{\text{mol}} - 6X$$

$$-2342 \frac{\text{kJ}}{\text{mol}} = -6X$$

$$X = 390 \text{ kJ/mol}$$

$x = \text{N-H Bond}$