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# Advanced Physical Chemistry Problems (V), Thermodynamics (ThermoChemistry)

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### Problems for the Advanced Physical Chemistry Student Part 5, Thermochemistry

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#### I. SYNOPSIS

This is a set of problems that were used near the turn of the century and which will be lost when the web site they were on disappears with my demise. Because these problems are being taken from the web and are being edited, their statements and the hints/answers offered are subject to the typical editorial errors that ensue when such work is undertaken in the vacuum of a non-teaching situation. Therefore, I claim any errors for myself, and hate to note that there most likely is no point in contacting me about them for obvious reasons.

#### II. THERMODYNAMICS, THE 2<sub>nd</sub> LAW

1. Calculate the expected adiabatic flame temperature of hydrogen burned in a stoichiometric amount of air. Assume  $(C_p(O_2)(g)) = 29.35 \text{ J/(mol^{\circ}K)}$ . Assume  $(C_p(H_2O(g)) = 33.577 \text{ and } \Delta H \text{ (standard,$  $formation)} = -241.814 \text{ kJ/mol}$ . Assume  $(C_p(N_2(g)))$  $= 29.12 \text{ J/(mol^{\circ}K)}$ . Assume that the actual 'combustion' takes place at 25°C. Assume air is 80% nitrogen and 20% oxygen. Therefore, the equation

$$\Delta H_{combustion} = \int_{273+25}^{273+T_{flame}} \left( 2C_p(N_2) + C_p(H_2O) \right) dT$$

where the leading sign is because we need to reverse the heat to make it go into the exiting gases.

As a side note, if the  $C'_p s$  are given as functions of temperature, we have an interesting nested integral problem to deal with; something to think about.

2.  $\Delta H_f^o$  of HCl(g) is -92.31 kJ/mole at 25°C. The heat capacities are given in the form:

$$C_p(T) = A + BT + CT$$

where T is in  ${}^{o}$ K and  $C_{p}$  is in Joules/(mole K).

Name	А	В	С
$H_2(g)$	29.07	$0.83x10^{-3}$	$2.01x10^{-6}$
$\operatorname{Cl}_2(g)$	31.72	$1.01x10^{-2}$	$4.04x10^{-6}$
$\mathrm{H}\mathrm{Cl}(\mathrm{g})$	28.16	$1.81x10^{-3}$	$1.55x10^{-6}$

Calculate the heat of formation of HCl (in kJ/mole) at standard pressure but 257.5°C.

#### Answer and/or Hint

When we need to work at temperatures other than "table" temperatures, and when we need to work at pressures other than "table" pressures, we need to know how to extend "table" values to the new conditions, hence this problem.

We know that

so

$$\left(\frac{\partial H}{\partial T}\right)_p = C_p$$

$$\left(\frac{\partial \Delta H}{\partial T}\right)_p = \Delta C_p$$

which means that

$$\int_{T_{table}}^{T_{desired}} \left(\frac{\partial \Delta H}{\partial T}\right)_p dT = \int_{T_{table}}^{T_{desired}} \Delta C_p dT$$

Answer and/or Hint

We're burning  $H_2$  in air, in a perfect (stoichiometric) amount of air. There's a little elementary chemistry knowledge required now. Air is  $4/5 N_2$ and  $1/5 O_2$  so if we assume one mole of  $H_2$  then since we have 1/2 mole of air, which contains 2 moles  $N_2$  and creates 1 moles of  $H_2O$  according to the equation

$$\mathrm{H}_2(g) + \frac{1}{2}\mathrm{O}_2(g) \to \mathrm{H}_2\mathrm{O}(g)$$

at  $25^{\circ}$ C.

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and we know all the  $C'_p s$ , so the r.h.s is simple. The l.h.s. is

$$\Delta H(T_{desired}) - \Delta H(T_{table}) = \int_{T_{table}}^{T_{desired}} \Delta C_p dT$$

3. Calculate the expected adiabatic flame temperature of hydrogen burned in air at  $25^{\circ}$ C, in which the ratio of hydrogen to air is 1:2 (by moles). Assume

Name	$C_p \frac{Joules}{mol^{\circ}C}$	
$C_p(\mathrm{H}_2(g))$	29.0	
$C_p(\mathcal{O}_2(g))$	29.4	
$C_p(\mathbf{N}_2(g))$	29.1	
$C_p(\mathrm{H}_2\mathrm{O}(g))$	33.6	

and  $\Delta H_f^o(\mathrm{H}_2\mathrm{O}(g)) = -242kJ/mol.$  Assume that the actual 'combustion' takes place at 25°C. Assume air is 80% nitrogen and 20% oxygen.

#### Answer and/or Hint

This is a fun problem, as are all flame temperature problems, in that we're able to calculate something "useful".

We're burning  $H_2$  in air, in a 1:2 mole ratio. There's a little elementary chemistry knowledge required now. Air is  $4/5 N_2$  and  $1/5 O_2$  so if we assume one mole of H)2 then we have 2 moles of air, which contains 8/5 moles  $N_2$  and 2/5 moles of  $O_2$ . Therefore the actual stoichiometry we're dealing with is one mole of  $H_2$  versus 2/5 moles of  $O_2$ .

$$\mathrm{H}_{2}(g) + \frac{1}{2}\mathrm{O}_{2}(g) \to \mathrm{H}_{2}\mathrm{O}(g)$$

at 25°C. Clearly, we're lacking in  $O_2$  which means that the exit gas will be a mixture of water vapor,  $H_2$  (left over) and  $N_2$  (carried through from beginning to end, but not involved in the combustion). We need to heat all these effluent gases using the  $\Delta H$  of combustion of the amount of H<sub>2</sub> actually consumed.

The details are left to the interested student.

4. Saturated solutions of ammonium chloride are often used in sports to reduce the swelling of a sprained ankle. Calculate the final temperature, if the following reaction

 $NH_4Cl(s) + 10H_2O(liq) \rightarrow NH_4Cl(aq, 10H_2O)$ 

takes place starting at  $25^{\circ}$ C.

 $\Delta H_f^o(\mathrm{NH}_4\mathrm{Cl}(aq, 10\mathrm{H}_2\mathrm{O})) = -71.567 \text{ kcal/mol}$  and that of  $\mathrm{NH}_4\mathrm{Cl}(\mathrm{s}) = -75.15 \text{ kcal/mol}$ . The specific heat of the solution is  $3.77 \text{ kJ/(kg^o\mathrm{K})}$ .

Answer and/or Hint

$$\frac{-71.567 - (-75.16)kcal/mol}{3.77\frac{1.987}{8.314} \times 0.2335 \ kg/mol} = \Delta t$$

where the heat of reaction is begin obtained from he athlete's hand, hence the sign. (Only partial units have been used.)

Don't forget to lower the temperature by  $\Delta t$  from  $25^{o}$ C.

#### III. EPILOGUE

After editing this material for weeks, and continuously finding errors, some small, some huge, I have to wrap it up and send this off. If, in the years 2008-2010 or so, you come across an error, and you e-mail me, I will try to have it corrected.

But since this material is written in LaTeX there is some doubt whether or not I'll have access to a Linux machine, and access to the digitial commons site. You can try; we'll see what happens, if anything. Thanks to all the students over the last 45 years who've taught me Physical Chemistry.