

## CH 223 Practice Problem Set #1

*This is a **practice problem set** and not the actual graded problem set that you will turn in for credit.  
Answers to each problem can be found at the end of this assignment.*

Covering: **Chapter Twelve and Chapter Guide One**

Important Tables and/or Constants: **Table of Thermodynamic Values** (found at the end of CH 223 Problem Set #1 or here: <http://mhchem.org/thermo>)

---

- Which substance has the higher entropy in each of the following pairs?
  - dry ice (solid  $\text{CO}_2$ ) at  $-78^\circ\text{C}$  or  $\text{CO}_2(\text{g})$  at  $0^\circ\text{C}$
  - liquid water at  $25^\circ\text{C}$  or liquid water at  $50^\circ\text{C}$
  - pure alumina,  $\text{Al}_2\text{O}_3(\text{s})$ , or ruby (ruby is  $\text{Al}_2\text{O}_3(\text{s})$  in which some of the  $\text{Al}^{3+}$  ions in the crystalline lattice are replaced with  $\text{Cr}^{3+}$  ions.)
  - one mole of  $\text{N}_2(\text{g})$  at 1 bar pressure or one mole of  $\text{N}_2(\text{g})$  at 10 bar pressure (both at 298 K)
- By comparing the formulas for each pair of compounds, decide which is expected to have the higher entropy. Assume all are at the same temperature.
  - $\text{O}_2(\text{g})$  or  $\text{CH}_3\text{OH}(\text{g})$  (two substances with the same molar mass)
  - $\text{HF}(\text{g})$ ,  $\text{HCl}(\text{g})$ , or  $\text{HBr}(\text{g})$
  - $\text{NH}_4\text{Cl}(\text{s})$  or  $\text{NH}_4\text{Cl}(\text{aq})$
  - $\text{HNO}_3(\text{g})$ ,  $\text{HNO}_3(\text{l})$ , or  $\text{HNO}_3(\text{aq})$
- Use  $S^\circ$  values to calculate the entropy change,  $\Delta S^\circ$ , for each of the following processes and comment on the sign of the change.
  - $\text{LiOH}(\text{s}) \rightarrow \text{LiOH}(\text{aq})$  (Note:  $S^\circ(\text{LiOH}(\text{aq})) = 91.6 \text{ J/molK}$ )
  - $\text{Na}(\text{g}) \rightarrow \text{Na}(\text{s})$
  - $\text{Br}_2(\text{l}) \rightarrow \text{Br}_2(\text{g})$
  - $\text{HCl}(\text{g}) \rightarrow \text{HCl}(\text{aq})$  (Note:  $S^\circ(\text{HCl}(\text{aq})) = 56.5 \text{ J/molK}$ )
- Calculate the standard molar entropy change of formation ( $\Delta S_f^\circ$ ) for each of the following compounds from the elements at  $25^\circ\text{C}$ .
  - $\text{HCl}(\text{g})$
  - $\text{Ca}(\text{OH})_2(\text{s})$
- Calculate the standard molar entropy change for each of the following reactions at  $25^\circ\text{C}$ . Comment on the sign of  $\Delta S^\circ$ .
  - $2 \text{Al}(\text{s}) + 3 \text{Cl}_2(\text{g}) \rightarrow 2 \text{AlCl}_3(\text{s})$
  - $2 \text{CH}_3\text{OH}(\text{l}) + 3 \text{O}_2(\text{g}) \rightarrow 2 \text{CO}_2(\text{g}) + 4 \text{H}_2\text{O}(\text{g})$
- Classify each of the reactions according to their spontaneity. Are these reactions enthalpy and/or entropy driven?
  - $\text{Fe}_2\text{O}_3(\text{s}) + 2 \text{Al}(\text{s}) \rightarrow 2 \text{Fe}(\text{s}) + \text{Al}_2\text{O}_3(\text{s})$   $\Delta H^\circ = -851.5 \text{ kJ}$   $\Delta S^\circ = -375.2 \text{ J/K}$
  - $\text{N}_2(\text{g}) + 2 \text{O}_2(\text{g}) \rightarrow 2 \text{NO}_2(\text{g})$   $\Delta H^\circ = 66.2 \text{ kJ}$ ;  $\Delta S^\circ = -121.6 \text{ J/K}$
- Heating some metal carbonates, among them calcium carbonate, leads to their decomposition.
$$\text{CaCO}_3(\text{s}) \rightarrow \text{CaO}(\text{s}) + \text{CO}_2(\text{g})$$
  - Calculate  $\Delta H^\circ$  and  $\Delta S^\circ$  for the reaction.
  - Is the reaction spontaneous at 298 K?
  - Is the reaction predicted to be spontaneous at higher temperatures?

8. Using values of  $\Delta H^\circ_f$  and  $S^\circ$ , calculate  $\Delta G^\circ_{\text{rxn}}$  for the following reaction. Is the reaction product-favored? Is the reaction enthalpy or entropy driven?
- $$2 \text{Pb(s)} + \text{O}_2(\text{g}) \rightarrow 2 \text{PbO(s, yellow)}$$
9. Using values of  $\Delta G^\circ_f$ , calculate  $\Delta G^\circ_{\text{rxn}}$  for each of the following reactions. Which are product-favored?
- $2 \text{K(s)} + \text{Cl}_2(\text{g}) \rightarrow 2 \text{KCl(s)}$
  - $2 \text{CuO(s)} \rightarrow 2 \text{Cu(s)} + \text{O}_2(\text{g})$
  - $4 \text{NH}_3(\text{g}) + 7 \text{O}_2(\text{g}) \rightarrow 4 \text{NO}_2(\text{g}) + 6 \text{H}_2\text{O(g)}$
10. For the reaction:  $\text{BaCO}_3(\text{s}) \rightarrow \text{BaO(s)} + \text{CO}_2(\text{g})$ ,  $\Delta G^\circ_{\text{rxn}} = +219.7 \text{ kJ}$ . Using this value and a table of thermodynamic data, calculate the value of  $\Delta G^\circ_f$  for  $\text{BaCO}_3(\text{s})$ .
11. Estimate the temperature required to decompose  $\text{HgS(s)}$  into  $\text{Hg(l)}$  and  $\text{S(g)}$ .

### Answers to the Practice Problem Set:

1. a.  $\text{CO}_2(\text{g})$  b.  $50^\circ \text{H}_2\text{O}(\text{l})$  c. ruby d.  $\text{N}_2(\text{g})$  at 1 bar
2. a.  $\text{CH}_3\text{OH}(\text{g})$  b.  $\text{HBr}(\text{g})$  c.  $\text{NH}_4\text{Cl}(\text{aq})$  d.  $\text{HNO}_3(\text{g})$
3. a. 48.8 J/K (entropy increase) b. -102.4 J/K (entropy decrease) c. 93.3 J/K (entropy increase) d. -130.4 J/K (entropy decrease) (*Instructor note: your values of  $\Delta S$  might be slightly different depending on the textbook used, etc., but they should be close to these values.*)
4. a. 10.0 J/K b. -294.1 J/K (see note in answer #17, above)
5. a. -504.6 J/K (entropy increase) b. 313.6 J/K (entropy decrease) (see note in answer #17, above)
6. a. enthalpy driven, spontaneous at low temperatures b. non-spontaneous at all temperatures
7. a.  $\Delta H^\circ = 191.59 \text{ kJ}$ ,  $\Delta S^\circ = 141.9 \text{ J/K}$  b. no c. yes
8.  $\Delta H^\circ = -434.64 \text{ kJ}$ ,  $\Delta S^\circ = -197.4 \text{ J/K}$ ,  $\Delta G^\circ = -375.77 \text{ kJ}$ ; product favored, enthalpy driven
9. a.  $\Delta G^\circ = -817.0 \text{ kJ}$ ; product favored b.  $\Delta G^\circ = 259.4 \text{ kJ}$ ; reactant favored c.  $\Delta G^\circ = -1101.3 \text{ kJ}$ ; product favored
10.  $\Delta G_f^\circ[\text{BaCO}_3(\text{s})] = -1134.4 \text{ kJ/mol}$
11. 2089 K or greater