Name	
Period	Date

20 • Entropy and Free Energy

- 1. Which of the following represents an increase in entropy?
 - a) freezing of water
 - b) boiling of water
 - c) crystallization of salt from a supersaturated solution
 - d) the reaction $2 \text{ NO}(g) \rightarrow N_2O_2(g)$
 - e) the reaction $2 H_2(g) + O_2(g) \rightarrow 2 H_2O(g)$
- The enthalpy of vaporization of methanol (CH₃OH) is 35.3 kJ/mol at the boiling point of 64.2 °C. Calculate the entropy change for methanol going from a liquid to vapor.
 - a) 600. J/K·mol d) -105 J/K·mol
 - b) 551 J/K·mol e) -551 J/K·mol
 - c) 105 J/K·mol
- 3. Calculate the standard entropy change for the following reaction,
 - $Cu(s) + \frac{1}{2}O_2(g) \rightarrow CuO(s)$

given that

$$S^{\circ}[Cu(s)] = 33.15 \text{ J/K} \cdot \text{mol}$$

$$S^{\circ}[O_{2}(g)] = 205.14 \text{ J/K} \cdot \text{mol}$$

$$S^{\circ}[CuO(s)] = 42.63 \text{ J/K} \cdot \text{mol}$$
a) 195.66 J/K d) -93.09 J/K
b) 02.00 J/K = 105.66 J/K

- b) 93.09 J/K e) 195.66 J/K
- c) -45.28 J/K
- 4. In which of the following reactions do you expect to have a decrease in entropy?
 - a) $Fe(s) \rightarrow Fe(l)$
 - b) $Fe(s) + S(s) \rightarrow FeS(s)$
 - c) 2 Fe(s) + 3/2 O₂(g) \rightarrow Fe₂O₃(s)
 - d) $HF(l) \rightarrow HF(g)$
 - e) $2 \operatorname{H}_2O_2(l) \rightarrow 2 \operatorname{H}_2O(l) + O_2(g)$

- PRACTICE TEST
- 5. The formation $\frac{1}{2} A_2 + 2 B_2 + C \rightarrow CAB_4$ has an enthalpy of formation of -104 kJ and a change in entropy of -60.8 J/K at 30 °C. What is ΔG and spontaneity of the reaction?
 - a) -85.6 kJ, spontaneous
 - b) -18.3 kJ, not spontaneous
 - c) +18.3 kJ, spontaneous
 - d) +85.6 kJ, not spontaneous
 - e) -85.6 kJ, not spontaneous
- 6. If ΔH and ΔS are both negative or positive,
 - then ΔG has a _____ sign.
 - a) positive d) large
 - b) negative e) no
 - c) variable
- 7. At what temperature would a given reaction become spontaneous if $\Delta H = +119$ kJ and $\Delta S = +263$ J/K?
 - a) 452 K d) 2.21 K
 - b) 2210 K e) 363 K
 - c) 382 K
- 8. The free energy change for a given reaction is -36.2 kJ. What is the equilibrium constant at 298 K?
 a) 0.985 d) 8.32 x 10⁻⁷
 - a) 0.985b) 2.22×10^6 c) 3.25×10^6 e) 3.25×10^6
 - c) 1.01

9. Given the following information, calculate ΔG° for the reaction below at 25°C: $SnCl_4(l) + 2 H_2O(l) \rightarrow SnO_2(s) + 4 HCl(g)$

- $\Delta H^{\circ} = 133.0 \text{ kJ} \text{ and } \Delta S^{\circ} = 401.5 \text{ J/K}$ a) -252.6 kJ d) 122.9 kJ
- b) -13.4 kJ e) 252.6 kJ
- c) 13.4 kJ

10. Given the following information, calculate ΔG° for the reaction below at 25°C: 2 H₂O₂(1) \rightarrow 2 H₂O(1) + O₂(g)

Compound	$\Delta H^{\circ}(kJ/mol)$	$\underline{S^{\circ}(J/K \cdot mol)}$
$H_2O_2(1)$	-187.8	109.6
$H_2O(l)$	-285.8	69.9
$O_2(g)$		205.1
a) -37700 l	kJ d)	-233.5 kJ
b) -342.6 k	J e)	-157.9 kJ
c) -233.5 k	J	

11. For the process at 25°C

	$I_2(g)$	$\rightarrow I_2(s)$	
what are	the signs	of ΔG , ΔH	, and ΔS ?

	<u>ΔG</u>	<u>ΔH</u>	<u>ΔS</u>
a)	+	_	_
b)	_	_	_
c)	_	+	+
d)	_	_	+
e)	+	+	+

- 12. If a process is exothermic and not spontaneous, then what must be true?
 - a) $\Delta S > 0$ d) $\Delta S < 0$
 - b) $\Delta H > 0$ e) $\Delta H = 0$
 - c) $\Delta G = 0$
- 13. For any reaction at equilibrium, which of the following is true?
 - a) $\Delta H < 0$ d) $\Delta H = 0$
 - b) $\Delta S = 0$ e) $\Delta G = 0$
 - c) $\Delta S < 0$
- 14. All of the following have $\Delta G^{\circ}_{f} = 0$ EXCEPT
 - a) $O_2(g)$ d) Ca(s)
 - b) $Br_2(g)$ e) Hg(l)
 - c) $H_2(g)$

Answers: 1B 2C 3D 4C 5A 6C 7A 8B 9C 10C 11B 12D 13E 14B 15B

- 15. Ammonium nitrate spontaneously dissolves in water at room temperature and the process causes the solution to become quite cold. Which of the following is TRUE about the dissolution of ammonium nitrate?
 - a) The process is exothermic.
 - b) Its solubility will be greater in warmer water.
 - c) ΔS° for the reaction is negative.
 - d) All solutions of ammonium nitrate are supersaturated.
 - e) All solutions of ammonium nitrate are cold.

Answers:



From the AP Exam formula sheet:

THERMOCHEMISTRY

$$\Delta S^{\circ} = \sum S^{\circ} \text{ products } -\sum S^{\circ} \text{ reactants}$$

$$\Delta H^{\circ} = \sum \Delta H_{f}^{\circ} \text{ products } -\sum \Delta H_{f}^{\circ} \text{ reactants}$$

$$\Delta G^{\circ} = \sum \Delta G_{f}^{\circ} \text{ products } -\sum \Delta G_{f}^{\circ} \text{ reactants}$$

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

$$= -RT \ln K = -2.303 RT \log K$$

$$= -n \mathcal{F} E^{\circ}$$

$$\Delta G = \Delta G^{\circ} + RT \ln Q = \Delta G^{\circ} + 2.303 RT \log Q$$

$$q = mc\Delta T$$

$$C_{p} = \frac{\Delta H}{\Delta T}$$