CHEM 1B - B.Kunkel

INSTRUCTIONS: Bubble in **Form A** on your Parscore Form

Write your perm number and **bubble in your perm number** (7 digits only)

Completely fill bubbles / answers





There are 20 questions. Each question is worth 5 points. It is to your benefit to **SHOW ALL YOUR WORK ON THE EXAM and CIRCLE YOUR ANSWERS. Turn in the Parscore form only**. Keep the exam so you can check your work and your answers later. The answers to the exam will be posted on our course web page. **NOTE:** No hats with caps allowed. No sharing of calculators. No NOTES allowed. **Cell phones and all other electronic devices must be turned off and put away.**

1. At constant pressure, the reaction is

$$2\mathrm{NO}_2(g) \to \mathrm{N}_2\mathrm{O}_4(g)$$

- A) spontaneous at low temperatures but not at high temperatures.
- B) spontaneous at high temperatures but not at low temperatures.
- C) always spontaneous.
- D) never spontaneous.
- 2. Calculate the standard heat of formation, ΔH_f° , for FeS₂(s), given the following information:

$$2\text{FeS}_2(s) + 5\text{O}_2(g) \rightarrow 2\text{FeO}(s) + 4\text{SO}_2(g)$$

$$\Delta H^{\circ}_{rxn} = -1370 \text{ kJ}$$

$$\Delta H_f^{\circ}$$
 for $SO_2(g) = -297 \text{ kJ/mol}$

$$\Delta H_f^{\circ}$$
 for FeO(s) = -268 kJ/mol

- A) +808 kJ
- B) -1550 kJ
- C) -774 kJ
- D) -686 kJ
- E) -177 kJ
- 3. The specific heat capacities of three metals are given below. If 1.00 g of each metal is heated to 100 °C and added to 10.0 g of H₂O at 25.0 °C, what is the order of the temperatures of the final mixtures from the lowest to the highest?

Metal	Specific heat, J/g°C
Fe	0.470
Pb	0.130
Zn	0.388

- A) Fe < Zn < Pb
- B) Pb < Zn < Fe
- C) Zn < Pb < Fe
- D) Zn < Fe < Pb
- E) none of the above

- 4. 2 mol of gas initially at 325 K are placed in a piston. The gas is cooled from 325 K to 225 K against a constant external pressure of 3 atm. Calculate the work.
- A) 16.4 J
- B) 2,494 J
- C) -1,663 J
- D) 1,663 J
- E) none of the above

- 5. At a constant pressure, the complete combustion of 1.00 mole of methane, CH₄, releases 802 kJ of heat. When 3.00 mol of oxygen gas react completely with excess methane at a constant pressure, what is the change in enthalpy?
- A) -1203 kJ
- B) -802 kJ
- C) -2406 kJ
- D) 1203 kJ
- E) none of the above

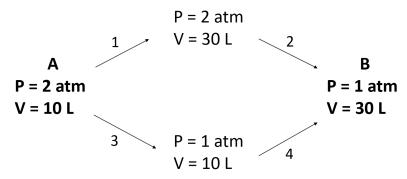
- 6. The following reaction has a ΔG° value of 42.6 kJ/mol at 25°C. Calculate K_a for the acid HB at 25 °C. HB(aq) + H₂O(l) \Longrightarrow H₃O⁺(aq) + B⁻(aq)
- A) 42,600
- B) 1.63
- C) 3.41×10^{-8}
- D) 14.0
- E) -17.2

- 7. For the reaction $2HF(g) \rightleftharpoons H_2(g) + F_2(g)$, $\Delta G^{\circ} = 28.3$ kJ at 1000 K. If, at this temperature, 5.00 mol of HF(g), 0.500 mol of $H_2(g)$, and 0.75 mol of $F_2(g)$ are mixed in a 1.00-L container, which of the following will happen?
- A) Some HF will be formed (from H_2 and F_2).
- B) The system is at equilibrium.
- C) Some HF will decompose (to yield H_2 and F_2).

- 8. The heat of vaporization for a compound is 17.2 kJ/mol at 25 °C. What is the change of entropy for one mole of the liquid compound when it vaporizes at 25 °C?
- A) 57.7 J/K
- B) 0.688 J/K
- C) $5.13 \times 10^3 \text{ kJ/K}$
- D) 3.16 J/K
- E) 239 J/K

- 9. Calculate the change in entropy when 2.0 mol of water are heated from 50 °C to 150 °C at P = 1 atm. $\Delta H_{fusion} = 6.01$ kJ/mol, $\Delta H_{vaporization} = 40.7$ kJ/mol, $C_{H2O(s)} = 2.01$ J/g °C, $C_{H2O(l)} = 4.18$ J/g °C, $C_{H2O(g)} = 2.03$ J/g °C
- A) 80.0 J/K
- B) 249 J/K
- C) 165 J/K
- D) 30.86 J/K
- E) none of the above

Questions 10 and 11. Consider 2.00 moles of a monatomic ideal gas that is taken from state A to state B by the following two different pathways:



- 10. Consider the above information. Calculate w from A to B via steps 3 and 4.
- A) -2030 J
- B) -20 J
- C) 0 J
- D) 3560 J
- E) 2500 J
- 11. Consider the above information. Calculate ΔE from A to B via steps 1 and 2.
- A) 2500 J
- B) 1,500 J
- C) 5500 J
- D) -2030 J
- E) 0 J
- 12. The standard enthalpy of reaction, ΔH° at 25 C° is -175.9 kJ mol⁻¹. Determine the value of $\Delta E_{\rm rxn}$ for this reaction at a constant pressure.

$$NH_3(g) + HCl(g) \rightarrow NH_4Cl(s)$$

- A) -164.8 kJ/mol
- B) -170.9 kJ/mol
- C) -175.9 kJ/mol
- D) -180.9 kJ/mol
- E) +5134 kJ/mol

13. In a well-insulated container, 50.0 g ice at 0.0 °C is added to 350. g water at 32.0 °C. What is the final temperature when the mixture reaches equilibrium? $\Delta H_{fusion} = 6.01 \text{ kJ/mol}$,

 $\Delta H_{\text{vaporization}} = 40.7 \text{ kJ/mol}, C_{\text{H2O(s)}} = 2.01 \text{ J/g} \,^{\circ}\text{C}, C_{\text{H2O (l)}} = 4.18 \text{ J/g} \,^{\circ}\text{C}, C_{\text{H2O(g)}} = 2.03 \text{ J/g} \,^{\circ}\text{C}$

- A) 18.0 °C
- B) 20.6 °C
- C) 22.0 °C
- D) 28.0 °C
- E) none of the above

- 14. A sample contains 2 mol of Cl_2 (g) ($C_p = 33.1$ J/mol K) is heated from 298 K to 350 K at constant pressure of 1 atm. Calculate ΔE .
- A) 3,442 J
- B) -865 J
- C) 1,209 J
- D) 2,577 J
- E) none of the above

15. Using the following reaction, how much heat is released when 10.0 g Fe and 2.00 g of oxygen gas are reacted at a constant pressure of 1 atm?

$$4\text{Fe (s)} + 3 \text{ O}_2 \text{ (g)} \rightarrow 2 \text{ Fe}_2\text{O}_3 \text{ (s)} \Delta H = -1652 \text{ kJ}$$

- A) 73.8 kJ
- B) 34.4 kJ
- C) 68.8 kJ
- D) 103 kJ
- E) none of the above

- 16. How much heat would your body lose if you were to ingest 800 g of ice at 0 °C and warm it to body temperature, 37 °C? Assume a constant pressure. $\Delta H_{fusion} = 6.01$ kJ/mol, $\Delta H_{vaporization} = 40.7$ kJ/mol, $C_{ice} = 2.01$ J/g °C, $C_{H2O(I)} = 4.18$ J/g °C, $C_{H2O(g)} = 2.03$ J/g °C
- A) 124 kJ
- B) 267 kJ
- C) 144 kJ
- D) 390 kJ
- E) none of the above

17. A coffee cup calorimeter having a heat capacity of 451 J/°C was used to measure the heat evolved when 0.0300 mol of NaOH(s) was added to 1000 mL of 0.0300 M HNO₃ initially at 23.000 °C. The temperature of the water rose to 23.639 °C. Calculate Δ H (in kJ/mol NaNO₃) for this reaction. Assume the specific heat of the final solution is 4.18 J/g°C; the density of each solution is 1.00 g/mL; and the addition of solid does not affect the volume of the solution.

$$HNO_3(aq) + NaOH(s) \rightarrow NaNO_3(aq) + H_2O(l)$$

- A) -63.7 kJ/mol
- B) -151 kJ/mol
- C) -2.55 kJ/mol
- D) -81.4 kJ/mol
- E) -98.6 kJ/mol

- 18. One mole of an ideal gas with a volume of 1.0 L and a pressure of 5.0 atm is allowed to expand isothermally into an evacuated bulb to give a total volume of 2.0 L. Calculate ΔS .
- A) 0 J
- B) -5.76 J
- C) 5.76 J
- D) 506 J
- E) none of the above

- 19. Substance X has a heat of vaporization of 55.4 kJ/mol at its normal boiling point (423°C). For the process $X(l) \to X(g)$ at 1 atm and 423°C, calculate the value of ΔS_{surr}
- A) -79.6 J/K•mol
- B) 0 J/K•mol
- C) -103 J/K•mol
- D) 103 J/K•mol
- E) 79.6 J/K•mol

- 20. The molar entropy of CH₄ (g) is 189 J/mol K at 25 $^{\circ}$ C and 1 atm pressure. Assuming ideal gas behavior, calculate the entropy of 0.39 mol CH₄ at 25 $^{\circ}$ C and a volume of 84 L.
- A) 196.1 J/K
- B) 80.8 J/K
- C) 207.1 J/K
- D) 73.7 J/K
- E) none of the above