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This test is closed note/book. One 8.5 x 11 handwritten crib sheet (one sided) is permitted.

Use a #2 pencil. Calculators are permitted. Computers, PDAs, and other electronic devices with a keyboard are not. Please turn off your cell phone. Cell phones may not be used as calculators.

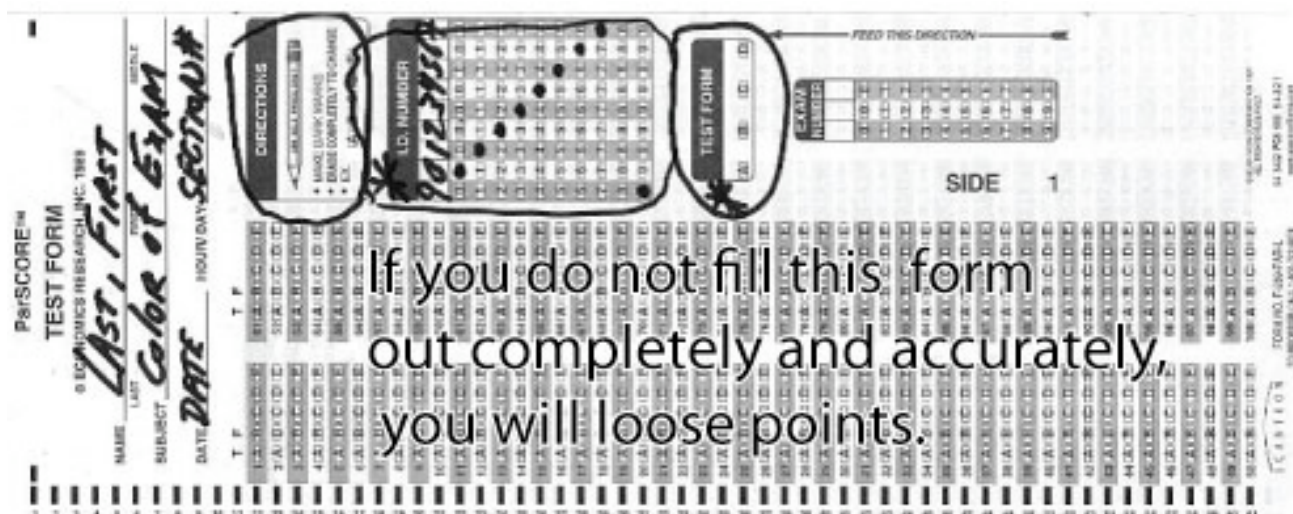
Write your name on this exam. Complete the Scantron card as shown below. You must bubble in your ID number, write in your section number and identify your Test Form (see top of this page). Scantron errors and omissions are punishable by point deductions.

When you take the exam, bubble in the scantron form *and* circle your answers on this exam. You must hand in both the scantron and the exam.

A total of 50 minutes is allotted for the exam. There are 20 questions. Each is worth five points. Answer every question. There is no penalty for guessing.

Circle Your Section Number

A1	M 2-3PM	CoC 52
A2	M 2-3PM	CoC 53
A3	M 2-3PM	MSE 1201A
A4	M 2-3PM	MSE 1222
A5	M 2-3PM	MSE 1224
B1	M 3-4PM	CoC 52
B2	M 3-4PM	CoC 53
B3	M 3-4PM	MSE 12101A
B4	M 3-4PM	MSE 1222
B5	M 3-4PM	MSE 1224



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1. You are given a solution of the weak base Novocain, Nvc. The pH of the solution is 11.0. You add the chloride salt of the conjugate acid of Novocain, NvcH^+Cl^- . Which statement is true?

- A) The pH and the pOH both increase.
 B) The pH and the pOH both decrease.
 C) The pH and the pOH remain unchanged.
 D) The pH increases and pOH decreases.
 E) The pH decreases and the pOH increases.

Answer: E

Chapter 7

2. Calculate $[\text{H}^+]$ in a solution that is 0.24 M in NaF and 0.60 M in HF ($K_a = 7.2 \times 10^{-4}$).

- A) 0.60 M
 B) 2.9×10^{-4} M
 C) 1.8×10^{-3} M
 D) 2.1×10^{-2} M
 E) 1.0×10^{-4} M

Answer: C

	HF	H^+	F^-
I	0.6	0	0.24
C	-x	x	x
E	0.60-x	x	0.24+x

$$K_a = \frac{x(0.24+x)}{(0.6-x)} = \frac{x(0.24)}{0.6} \text{ (approx)}$$

$$(7.2 \times 10^{-4})0.6/(0.24) = 1.8 \times 10^{-3}$$

Chapter 7

3. For ammonia (NH_3), K_b is 1.8×10^{-5} . The buffering capacity of a 1 M solution of NH_4Cl is at a maximum at a pH of

- A) 4.7
 B) 7.2
 C) 12.2
 D) 9.3
 E) none of these

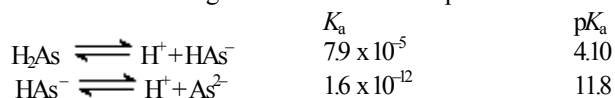
Answer: D

$$\text{p}K_b = -4.75$$

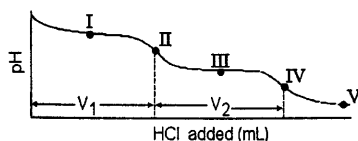
$\text{p}K_a = 9.3$, buffering capacity is max when $\text{pH} = \text{p}K_a$

Chapter 8

4. Consider the following information about diprotic ascorbic acid (H_2As).



The curve for titration of disodium ascorbate (Na_2As) with HCl is shown below:



What major species is (are) present at point III of the titration curve (note that the titration starts with essentially 100% As^{2-})?

- A) As^{2-} and HAS^-

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- B) HAs^- only
- C) HAs^- and H_2As
- D) H_2As only
- E) H_2As and H^+

Answer: C
Chapter 8

5. Referring to the disodium ascorbate titration above: which of the following is a major species present at point IV?

- A) H_2As
- B) HAs^-
- C) As^{2-}
- D) H^+
- E) none of these

Answer: A
Chapter 8

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6. A gas releases 3.8 J of heat to the surroundings and performs 13.7 J of work. What is the change in energy of the gas?

- A) -17.6 J
- B) 17.6 J
- C) -9.9 J
- D) 9.9 J
- E) 3.8 J

Answer: A

$$\Delta E = q + w = -3.8 \text{ J} + (-13.7 \text{ J}) = -17.6 \text{ J}$$

Chapter 9

6. (question not used) Calculate ΔE for a system that releases 35 J of heat to the surroundings while 54 J of work is done on it.

- A) -89 J
- B) -19 J
- C) 19 J
- D) 89 J
- E) 35 J

Answer: C

$$\Delta E = q + w = -35 \text{ J} + 54 \text{ J} = +19 \text{ J}$$

Chapter 9

7. Which of the following statements is false?

- A) In going from a particular set of reactants to a particular set of products, the change in enthalpy is the same, whether the reaction takes place in a series of steps or in a single step.
- B) The entropy of the universe increases for any spontaneous process.
- C) The energy of the universe is conserved.
- D) A system will always proceed spontaneously to the arrangement with the lowest enthalpy.
- E) Energy cannot be created or destroyed.

Answer: D

7 (question not used). Calculate the work for the expansion of an ideal gas from 3.0 to 6.0 L against an external pressure of 1.6 atm at constant temperature.

- A) 4.8 L·atm
- B) -4.8 L·atm
- C) 0.0 L·atm
- D) 5.6 L·atm
- E) -1.9 L·atm

Answer: B

$$w = -P\Delta V = -1.6 \text{ atm} (6.0 \text{ L} - 3.0 \text{ L}) = -4.8 \text{ L}\cdot\text{atm}$$

Chapter 9

7 (question not used). Calculate the work for the compression of an ideal gas from 6.0 to 3.0 L against an external pressure of 1.6 atm at constant temperature.

- A) 4.8 L·atm
- B) -4.8 L·atm
- C) 0.0 L·atm
- D) 5.6 L·atm
- E) -1.9 L·atm

Answer: A

$$w = -P\Delta V = -1.6 \text{ atm} (3.0 \text{ L} - 6.0 \text{ L}) = 4.8 \text{ L}\cdot\text{atm}$$

Chapter 9

Consider the following process carried out on 1.0 mol of a monatomic ideal gas at constant pressure:

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Start: (3.00 atm, 20.0 L)
End: (3.00 atm, 50.0 L).

8. Calculate the work, w .

- A) -90 L•atm
- B) 90 L•atm
- C) -30 L•atm
- D) 30 L•atm
- E) 0 L•atm

Answer: A

$$w = -P\Delta V = -3 \text{ atm} (50 \text{ L} - 20 \text{ L}) = -90 \text{ L}\cdot\text{atm}$$

Chapter 9

9. Calculate the heat, q .

- A) 226 L•atm
- B) -226 L•atm
- C) 135 L•atm
- D) -135 L•atm
- E) none of these

Answer: A

$$T(A) = PV/nR = (3 \text{ atm})(20 \text{ L})/[(1.0 \text{ mol})(0.082 \text{ L}\cdot\text{atm mol}^{-1} \text{ K}^{-1})] = 731 \text{ K}$$

$$T(B) = PV/nR = (3 \text{ atm})(50 \text{ L})/[(1.0 \text{ mol})(0.082 \text{ L}\cdot\text{atm mol}^{-1} \text{ K}^{-1})] = 1830 \text{ K}$$

$$\Delta T = 1099 \text{ K}$$

$$q = nC_p\Delta T = (1 \text{ mol})(5/2 R \text{ J mol}^{-1} \text{ K}^{-1})(101 \text{ J/L}\cdot\text{atm})^{-1}(1099 \text{ K}) \\ = (5/2 \times 8.31) 1099/101 = 226 \text{ L}\cdot\text{atm}$$

10. The same amount of heat is added to two metal blocks of equal number of moles but made of different metals. Which undergoes the smaller change in temperature?

- A) The metal with the higher heat capacity.
- B) The metal with the lower heat capacity.
- C) Both undergo the same change in temperature.
- D) To determine this, you need to know the initial temperatures of the metals.
- E) To determine this, you need to know which metals you are talking about.

Answer: A

Chapter 9

11. As a warm brick (the system) spontaneously cools in a cold pool of water (the surroundings), the entropy of the brick (S_{system})

- A) increases
- B) decreases
- C) does not change
- D) changes in a way that cannot be predicted
- E) all of the above

Answer: B

Chapter 10

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12. As the warm brick (the system) spontaneously cools in the cold pool of water (the surroundings), the entropy of the surroundings ($S_{\text{surroundings}}$)

- A) increases
- B) decreases
- C) does not change
- D) changes in a way that cannot be predicted
- E) all of the above

Chapter 10

Answer: A

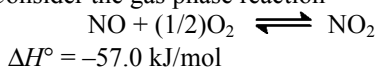
13. As the warm brick (the system) spontaneously cools in the cold pool of water (the surroundings),

- A) $|\Delta S_{\text{system}}| > |\Delta S_{\text{surroundings}}|$
- B) $|\Delta S_{\text{system}}| < |\Delta S_{\text{surroundings}}|$
- C) $|\Delta S_{\text{system}}| = |\Delta S_{\text{surroundings}}|$
- D) the relationship of $|\Delta S_{\text{system}}|$ and $|\Delta S_{\text{surroundings}}|$ cannot be predicted
- E) none of the above

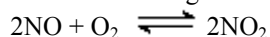
Chapter 10

Answer: B

14. Consider the gas phase reaction



What is ΔH° for the following reaction:



- A) 57.0 kJ
- B) -114 kJ
- C) 114 kJ
- D) -28.5 kJ
- E) 778 kJ

Answer: B

Chapter 9

15. In a reaction where a diatomic molecule (for example O_2) spontaneously forms from its atoms at standard temperature and pressure, what are the signs of ΔH , ΔS , and ΔG , respectively?

- A) - - -
- B) + + +
- C) + - -
- D) - + +
- E) - - +

Answer: A

Chapter 10

16. Consider the spontaneous freezing of liquid water at -10°C . For this process what are the signs for ΔH , ΔS , and ΔG , respectively?

- A) + - 0
- B) - + 0
- C) - + -
- D) + - -
- E) - - -

Answer: E

Chapter 10

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17. Which one of the following processes has $\Delta H < 0$?

- A) freezing of liquid Hg
- B) combustion of cellulose
- C) cooling water
- D) all of the above (a-c)
- E) none of the above (a-c)

Chapter 9

Answer: D

18. In SI units the universal gas constant R is $8.31 \text{ J mol}^{-1}\text{K}^{-1}$. R is also $0.0820 \text{ L atm mol}^{-1}\text{K}^{-1}$. Therefore, 1.00 L-atm is equivalent to how many J?

- A) 9.87
- B) 9.87×10^{-3}
- C) 101.3
- D) 1.013×10^5
- E) none of these are correct.

Answer: C

19. For a balloon expanding against atmospheric pressure, the work done on the air in the balloon is

- A) positive
- B) negative
- C) of indeterminate sign
- D) zero
- E) this cannot be answered without additional information.

Answer: B

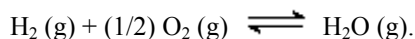
20. This is Test Form (look at the top of the page):

- A) A
- B) B
- C) C
- D) D

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Extra Credit / Bonus: The following bonus questions are worth just one point (1 pt) each!

Consider the gas-phase reaction



Standard gas phase thermodynamic information is available:

$$\begin{array}{ll} \Delta H_f^\circ(\text{H}_2) = 0.0 & S_f^\circ(\text{H}_2) = 131 \text{ J/mol-K} \\ \Delta H_f^\circ(\text{O}_2) = 0.0 & S_f^\circ(\text{O}_2) = 205 \text{ J/mol-K} \\ \Delta H_f^\circ(\text{H}_2\text{O}) = -242 \text{ kJ/mol} & S_f^\circ(\text{H}_2\text{O}) = 189 \text{ J/mol-K} \end{array}$$

21. What is ΔH_r° for this reaction?

- A) + 121 kJ/mol B) - 121 kJ/mol C) + 242 kJ/mol D) - 242 kJ/mol
E) Cannot be determined from the information given.

Answer: D

22. What is ΔS_r° for this reaction?

- A) + 44 J/mol-K B) - 44 J/mol-K C) + 147 J/mol-K D) -147 J/mol-K
E) Cannot be determined from the information given.

Answer: B

$$189 - (1/2)205 - 131 = -44$$

23. What is ΔG_r° for this reaction at 298K?

- A) -255 kJ/mol B) -229 kJ/mol C) +229 kJ/mol D) +255 kJ/mol
E) Cannot be determined using the information given.

Answer: B

$$-242 - (298)(-44/1000) = -229$$

24. For this reaction, at about what temperature is $\Delta G_r^\circ = 0$?

- A) 5.5 K B) 300 K C) 5200 K D) 5500 K
E) Cannot be determined using the information given.

Answer: D

$$T = 242 / .044 = 5500 \text{ K}$$

25. For this reaction, what is the approximate value of $\ln(K_{\text{eq}})$ at $T = 1000 \text{ K}$?

- A) 24 B) 29 C) 34 D) -24
E) Cannot be determined using the information given.

Answer: A

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ = -242 - 1000(-0.044) = -198 = -RT \ln(K_{\text{eq}})$$

$$\ln(K_{\text{eq}}) = (198000 \text{ J/mol}) / [(8.31 \text{ J/mol K})(1000 \text{ K})] = 23.8$$

26. The maximum work (in absolute value) obtainable from this reaction at 1000 K, and $Q = 0.0025$ (Q is the reaction quotient) is approximately:

- A) 0 kJ/mol B) 150 kJ/mol C) 200 kJ/mol D) 250 kJ/mol
E) Cannot be determined using the information given.

Answer: D

$$w = -198000 + 8.3 \cdot 1000 \ln(0.0025) = -198000 \text{ J/mol} - 50000 \text{ J/mol} = -248 \text{ kJ/mol}$$

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1 mole atoms = 6.022×10^{23} atoms

$h = 6.626 \times 10^{-34} \text{ Js}$ 1 J (Joule) = $1 \text{ kg} \frac{\text{m}^2}{\text{s}^2}$

1 J (Joule) = .00987 L - atm (or 101.3 J/L - atm)

$c = 3.0 \times 10^8 \text{ m/s}$

Mass Electron = $9.10939 \times 10^{-31} \text{ kg}$

Mass Proton = $1.67 \times 10^{-27} \text{ kg}$

Mass Neutron = $1.67 \times 10^{-27} \text{ kg}$

$R = 0.0821 \frac{\text{L} \cdot \text{atm}}{\text{mol} \cdot \text{K}}$ $R = 8.31 \frac{\text{J}}{\text{mol} \cdot \text{K}}$

Vapor Pressure ($\text{H}_2\text{O}, 373\text{K}$) = 760 torr

Formal Charge = $V - (L + 0.5 S)$

V = Number of Valence Electrons

L = Number of Lone Pair Electrons

S = Number of Shared Electrons



$P = X_1 \cdot P^\circ$

$m = \text{molality} = \frac{\text{mol of solute}}{\text{kg of solvent}}$

$\Delta T_f = -m \cdot K_f$ $\Delta T_b = m \cdot K_b$

$X_1 = \text{mole fraction} = \frac{n_1}{n_{\text{total}}}$

$\Pi = MRT$

$q = mC\Delta T$ $q_v = nC_v\Delta T$

$q_p = nC_p\Delta T$ $dS = \frac{dq}{T}$

$\Delta G = \Delta H - T\Delta S$ $\Delta G^\circ = -RT \ln K$ $\Delta G = \Delta G^\circ + RT \ln Q$

$T_c = \frac{\Delta H}{\Delta S}$ $C_{\text{water}} = 4.184 \frac{\text{J}}{\text{gram K}}$

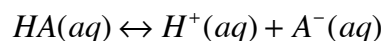
$c = \lambda\nu$ $E = mc^2$ $\lambda = \frac{h}{p}$

$\hat{H}\psi = E\psi$ $\Delta x \cdot m\Delta v \geq \frac{h}{4\pi}$

Maximum Occupancy = $2n^2$

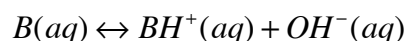
$pH = -\log[H^+]$ $pOH = -\log[OH^-]$

$K_w = K_a K_b$ $pH + pOH = 14$



$K_a = \frac{[H^+][A^-]}{[HA]}$

$K_w = 1 \times 10^{-14}$ at 25°C



$K_b = \frac{[BH^+][OH^-]}{[B]}$

$pK_a = -\log K_a$ $pK_w = 14$ at 25°C

$\Delta E = q + w$

$w = -P_{\text{ext}}\Delta V$

$C_p = \frac{5}{2}R$ (monoatomic ideal gas)

$C_v = \frac{3}{2}R$ (monoatomic ideal gas)

$\Delta H = q_p$ (constant temperature)

$w_{\text{rev}} = -nRT \ln\left(\frac{V_2}{V_1}\right)$

E(monatomic) = KE(mol) = $\frac{3}{2}(RT) = \frac{3}{2}(PV)$

KE(particle) = $\frac{1}{2}mu^2$

Print Name

The Periodic Table of the Elements

1 H Hydrogen 1.00794	4 Be Beryllium 9.012182	21 Sc Scandium 44.955910	22 Ti Titanium 47.867	23 V Vanadium 50.9415	24 Cr Chromium 51.9961	25 Mn Manganese 54.938049	26 Fe Iron 55.845	27 Co Cobalt 58.933200	28 Ni Nickel 58.6934	29 Cu Copper 63.546	30 Zn Zinc 65.39	31 Ga Gallium 69.723	32 Ge Germanium 72.61	33 As Arsenic 74.92160	34 Se Selenium 78.96	35 Br Bromine 79.904	36 Kr Krypton 83.80	2 He Helium 4.003				
3 Li Lithium 6.941	12 Mg Magnesium 24.3050	39 Y Yttrium 88.90585	40 Zr Zirconium 91.224	41 Nb Niobium 92.90638	42 Mo Molybdenum 95.94	43 Tc Technetium (98)	44 Ru Ruthenium 101.07	45 Rh Rhodium 102.90550	46 Pd Palladium 106.42	47 Ag Silver 107.8682	48 Cd Cadmium 112.411	49 In Indium 114.818	50 Sn Tin 118.710	51 Sb Antimony 121.760	52 Te Tellurium 127.60	53 I Iodine 126.90447	54 Xe Xenon 131.29					
19 K Potassium 39.0983	20 Ca Calcium 40.078	37 Rb Rubidium 85.4678	38 Sr Strontium 87.62	55 Cs Cesium 132.90545	56 Ba Barium 137.327	57 La Lanthanum 138.9055	72 Hf Hafnium 178.49	73 Ta Tantalum 180.9479	74 W Tungsten 183.84	75 Re Rhenium 186.207	76 Os Osmium 190.23	77 Ir Iridium 192.217	78 Pt Platinum 195.078	79 Au Gold 196.96655	80 Hg Mercury 200.59	81 Tl Thallium 204.3833	82 Pb Lead 207.2	83 Bi Bismuth 208.98038	84 Po Polonium (209)	85 At Astatine (210)	86 Rn Radon (222)	
87 Fr Francium (223)	88 Ra Radium (226)	89 Ac Actinium (227)	104 Rf Rutherfordium (261)	105 Db Dubnium (262)	106 Sg Seaborgium (263)	107 Bh Bohrium (262)	108 Hs Hassium (265)	109 Mt Meitnerium (266)	110 Ds Darmstadtium (269)	111 Rg Roentgenium (272)	112 Cn Copernicium (277)	113 Nh Nihonium (277)	114 Fl Flerovium (277)									
58 Ce Cerium 140.116	59 Pr Praseodymium 140.90765	60 Nd Neodymium 144.24	61 Pm Promethium (145)	62 Sm Samarium 150.36	63 Eu Europium 151.964	64 Gd Gadolinium 157.25	65 Tb Terbium 158.92534	66 Dy Dysprosium 162.50	67 Ho Holmium 164.93032	68 Er Erbium 167.26	69 Tm Thulium 168.93421	70 Yb Ytterbium 173.04	71 Lu Lutetium 174.967									
90 Th Thorium 232.0381	91 Pa Protactinium 231.03588	92 U Uranium 238.0289	93 Np Neptunium (237)	94 Pu Plutonium (244)	95 Am Americium (243)	96 Cm Curium (247)	97 Bk Berkelium (247)	98 Cf Californium (251)	99 Es Einsteinium (252)	100 Fm Fermium (257)	101 Md Mendelevium (258)	102 No Nobelium (259)	103 Lr Lawrencium (262)									

Answer: B

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