# THE UNIVERSITY OF CALGARY FACULTY OF SCIENCE CHEMISTRY 209

### Winter 2013

#### **Final Exam**

Date: April 19<sup>th</sup>, 2013 Time Allotted: 3 hours

<u>Please make sure you fill in your NAME, student ID and Lecture Section and encode all information on the optical score sheet.</u>

Failure to do so will result in the loss of at least 2 marks.

Only Casio FX260 or similar non-programmable scientific calculators are permitted.

A Chemical Data and Formula Sheet is attached, which can be detached.

FIRST NAM	Œ:		LAST I	LAST NAME:						
ID:										
Lecture Sect	ion: Please c	heck one								
	L01 L02	TR TR	12:30 pm 8:00 am	Dr. Masood Parvez Dr. Scott Hinman						

This exam consists of 14 pages, 21 multiple choice questions worth two marks each (total 42 marks), and 4 written answer questions (total 30 marks) for a total of 72 marks. Make sure to count the number of pages (should be 14 pages excluding data sheet) and LET THE INVIGILATOR KNOW IMMEDIATELY IF YOU ARE MISSING A PAGE.

All questions must be answered to earn full marks. The answers to the multiple choice questions must be entered on the optical score sheet using pencil. Answers to the written answer questions should be written in the exam booklet in ink.

## <u>AT THE END OF THE EXAMINATION, HAND IN THE WHOLE EXAM BOOKLET AND THE</u> <u>OPTICAL SCORE SHEET</u>

Do not write in spaces in the table below

Q 22	Q 23	Q24	Q25

### SECTION I - Machine graded section (Total value 42 marks) Select the best answer and encode it on the Optical Score Sheet

1. What is the mass percent of each element in sulfuric acid, H<sub>2</sub>SO<sub>4</sub>?

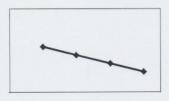
Molar mass of 
$$H_25O_4 = 98.086$$
  
%  $H = \frac{2.016}{98.086} \times 100 = 2.055\%$   
%  $S = \frac{32.07}{98.086} \times 100 = 32.70\%$ 

(a.) 2.055% H, 32.70% S, 65.25% O

b. 1.028% H, 32.69% S, 66.28% Oc. 28.57% H, 14.29% S, 57.17% O

$$\%0 = \frac{64.00}{98.086} \times 100 = 65.25\%$$

2. A student analyzed a first-order reaction and obtained the graph below. Unfortunately, the student forgot to label the axes. What are the correct labels for the *x* and *y* axes?



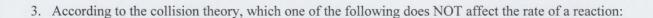
(a) 
$$x$$
 axis = time,  $y$  axis =  $ln[A]$ 

b. 
$$x \text{ axis} = \ln[\text{time}], y \text{ axis} = [A]$$

c. 
$$x = \ln[time], y = A$$

d. 
$$x$$
 axis = time,  $y$  axis =  $1/[A]$ 

e. 
$$x$$
 axis =  $1$ /time,  $y$  axis =  $1$ /[A]



4. Nitrogen dioxide reacts with carbon monoxide to produce nitrogen monoxide and carbon dioxide.

$$NO_2(g) + CO(g) \rightarrow NO(g) + CO_2(g)$$

A proposed mechanism for this reaction is:

$$2 \text{ NO}_2(g) \stackrel{\mathbf{k_1}}{\rightleftharpoons} \text{NO}_3(g) + \text{NO}(g)$$

$$\text{NO}_3(g) + \text{CO}(g) \stackrel{\mathbf{k_2}}{\rightarrow} \text{NO}_2(g) + \text{CO}_2(g)$$

(fast, equilibrium)

$$NO_3(g) + CO(g) \xrightarrow{K_2} NO_2(g) + CO_2(g)$$

(slow)

What is a rate law that is consistent with the proposed mechanism?

(a) rate = 
$$k [NO_2]^2 [CO] [NO]^{-1}$$

b. rate = 
$$k [NO_2]^2 [CO]$$

c. rate = 
$$k$$
 [NO<sub>2</sub>] [CO]

d. rate = 
$$k [NO_3] [CO]^2$$

e. rate = 
$$k [NO_2]^2$$

$$K_{1}[NO_{2}]^{2} = K_{-1}[NO_{3}][NO]$$

$$[NO_3] = \frac{k_1}{k_{-1}} \frac{[NO_2]^2}{[NO]}$$

Rate = 
$$k_2 \cdot \frac{k_1}{k_1} \cdot \frac{[NO_2]^2}{[NO]}$$
 (CO) or  $k[NO_2]$  [CO]  $[NO]$ 

At 25 °C, the equilibrium constant for the autoionization of water is  $K_W = 1.0 \times 10^{-14}$ . At 50 °C,

 $K_W = 5.5 \times 10^{-14}$ . Determine  $\Delta H$  for the autoionization of water.

$$\ln \left( \frac{5.5 \times 10^{-14}}{1.0 \times 10^{-14}} \right) = \frac{\Delta H}{8.314 \text{ J mol}^{-1} \text{ K}^{-1}} \left( \frac{1}{298} - \frac{1}{323} \right) \text{ K}$$

$$N_2 + 3H_2 \rightleftharpoons 2NH_3$$

 $N_2 + 3H_2 \implies 2NH_3$  is  $K_c = 6.0 \times 10^{-2}$ . The numerical

value of K<sub>p</sub> at this temperature is:

$$k_p = k_c (RT)^{\Delta n}$$

$$=6.0\times10^{-2}(0.08205\times773)^{-2}$$

$$= 6.0 \times 10^{-2} (63.42)^{-2} = 1.5 \times 10^{-5}$$

An evacuated 2.0 L flask was filled with 0.10 moles SO<sub>3</sub>(g) and 0.20 moles of NO (g). The following 7. reaction was allowed to come to equilibrium:  $SO_3(g) + NO(g) \Longrightarrow SO_2(g) + NO_2(g)$ If  $K_c$  for this reaction is 0.50, the equilibrium concentration of  $SO_2(g)$  is:

a. 
$$0.056 \,\mathrm{M}$$
  $0.05 - \infty$   $0.10 - \infty$   $0.05 - \infty$   $0.10 - \infty$ 

- 0.021 M b.
- 0.047 M c.
- 0.035 M d.
- (e.) 0.028 M

$$K_c = \frac{(x)(x)}{(0.05-x)(0.10-x)} = 0.50$$

Colgate PreviDent™ toothpaste contains 1.0 mass percent NaF. Given that HF is a weak acid with Ka = 8. 6.8 x 10<sup>-4</sup>, what mass of PreviDent™ toothpaste would have to be dissolved in pure water to give a total volume of 1.00 L of a solution with pH = 7.9? 0H = 6.1 :  $OH = 7.94328 \times 10^{-1}$ 

$$F + H_2O = HF + OH - Colgate PreviDent$$

$$210 g$$

$$K_b = \frac{1.0 \times 10^{-14}}{6.8 \times 10^{-4}} = \frac{(\pi)^2}{F - \pi}$$
:.  $F = 4.292268 \times 10^{-2}$  moles

2.0 g

a.

- In which of the following solutions would you expect LiF to be the most soluble given that Ksp for LiF 9. is  $1.7 \times 10^{-3}$ ?
- 100.0 mL of pure water a.
- 100.0 mL of 1.0 M HCl (b.)
- 100.0 mL of 1.0 M CH<sub>3</sub>CO<sub>2</sub>H c.
- d. 100.0 mL 0f 1.0 M NH3
- 100.0 mL of 1.0 M NaOH e.
- What is the resulting pH when 50.00 mL of 0.10 M HCl is added to a buffer prepared by combining 10. 0.100 moles of acetic acid and 0.110 moles of sodium acetate with 500 mL of pure water?  $K_a$  for acetic acid is  $1.8 \times 10^{-5}$ .

acid is 
$$1.8 \times 10^{-5}$$
.

Moles of HCl added =  $5.0 \times 10^{-3}$  moles

ACOH

0.110

0.100

-5.0×10<sup>3</sup>

+ \$6.5.0×10<sup>3</sup>

0.100+5.0×10

0.55

= 0.191 M

= 0.191 M

c. 
$$4.60$$
 (C)  $-5.0 \times 10$   $+645.0 \times 10$  d.  $5.12$  (E)  $0.110-5.0 \times 10^3$   $0.100+5.0 \times 10^3$  e.  $4.42$  (E)  $0.55$ 

$$pH = pk_{\alpha} + log \frac{[AcO-]}{[AcOH]}$$
 :  $pH = pk_{\alpha} = 4.74$ 

 $CaSO_4$  has  $K_{sp} = 2.4 \times 10^{-5}$  while  $BaSO_4$  has  $K_{sp} = 1.1 \times 10^{-10}$ . Determine the equilibrium constant,  $CaSO_4(s) + Ba^{2+}(aq) \rightleftharpoons Ca^{2+}(aq) + BaSO_4(s)$ K<sub>c</sub>, for the reaction:

b. 
$$2.4 \times 10^{-5}$$
  $Ba^{2+}(aq) + So_{4}(aq) = BaSo_{4}(s) / 1.1 \times 10^{-10}$ 

c. 
$$1.1 \times 10^{-10}$$
  $Da(aq) + SO_4(aq) = BaSO_4(s) / 1.1 \times 10^{-10}$ 

d. 
$$4.4 \times 10^{-6}$$
  
e.  $2.64 \times 10^{-15}$   $CaSO_4(5) + Ba^2(aq) = Ca^2(aq) + BaSO_4(5)$ 

$$= 2.4 \times 10^{-5} / 1.1 \times 10^{-10}$$
$$= 2.18 \times 10^{5}$$

#### Standard potentials at 25 °C which may be used in Questions 13 to 15 and 24

$$IO_4^- + 2H^+ + 2e^- \rightleftharpoons IO_3^- + H_2O$$
  $E^\circ = 1.589 \text{ V}$   
 $MnO_4^- + 8H^+ + 4e^- \rightleftharpoons Mn^{2+} + 4H_2O$   $E^\circ = 1.507 \text{ V}$   
 $Cl_2(g) + 2e^- \rightleftharpoons 2Cl^-(aq)$   $E^\circ = 1.360 \text{ V}$   
 $Fe^{3+} + 1e^- \rightleftharpoons Fe^{2+}$   $E^\circ = 0.771 \text{ V}$   
 $2H^+ + 2e^- \rightleftharpoons H_2(g)$   $E^\circ = 0.000 \text{ V}$   
 $HCO_2H + 2H^+ + 2e^- \rightleftharpoons H_2CO + H_2O$   $E^\circ = -0.029 \text{ V}$   
 $Pb^{2+} + 2e^- \rightleftharpoons Pb(s)$   $E^\circ = -0.126 \text{ V}$ 

- 13. Which species is the strongest oxidizing agent under standard conditions?
- Pb(s) a.
- (b.) IO4-
- $Pb^{2+}$
- d. IO3-
- H<sub>2</sub>O e.

Determine the equilibrium constant for the reaction between IO<sub>4</sub>- and Fe<sup>2+</sup>. 14.

c. 
$$0.57 \times 10^{13}$$

$$E' = \frac{0.0592}{2} \log K$$

$$log K = 27.635$$
 ...  $K = 4.32 \times 10^{27}$ 

15. What is the potential of the following cell?

Pb | Pb $^{2+}$ (0.200 M) || Cl $^{-}$ (0.200M) | Cl $_{2}$ (g, 0.400 atm.) | Pt

$$E = E - \frac{0.0592 \log Q}{2} \log Q = \frac{(0.200)(0.200)^2}{(0.400)}$$

19. An element with the outermost electron configuration

$$\begin{array}{ccc}
\uparrow\downarrow & & \uparrow\downarrow \uparrow\uparrow \uparrow\\ ns & & np
\end{array}$$

could be\_\_\_\_\_.

- 82Pb
- 40Zr b.
- 42Mo
- (d.) 34Se
- 19K

 $1s^2 2s^2 2p^6 3s^3 3p^6 3d^{10} 4s^2 4p^4$ 



#### SECTION II: Fill-in-the-blanks and long answer questions. (Total value 30 marks) Write your answers in ink in the space provided.

#### 22. Fill-in-the-blanks.

10 marks

Nitroglycerine decomposes violently according to the chemical equation below. What mass of carbon a. dioxide gas is produced from the decomposition of 5.00 g C<sub>3</sub>H<sub>5</sub>(NO<sub>3</sub>)<sub>3</sub>?

$$4 C_3H_5(NO_3)_3(\ell) \rightarrow 12 CO_2(g) + 6 N_2(g) + 10 H_2O(g) + O_2(g)$$
  
(5.00 / 255.084) (12 / 4) (44.1) = 2.93 g

2.93 g

If 5.15 g Fe(NO<sub>3</sub>)<sub>3</sub> is dissolved in enough water to make exactly 150.0 mL of solution, what is the molar b. concentration of nitrate ion?

$$(5.15 / 241.88) (3) (1 / 0.150) = 0.426 M$$

0.426 M

The rate constant at 366 K for a first-order reaction is  $7.7 \times 10^{-3} \text{ s}^{-1}$  and the activation energy is 15.9 c. kJ/mol. What is the value of the frequency factor, A, in the Arrhenius equation?

$$K = A e^{-15.9/366 \times 8.314 \times 10_{.3}}$$

$$7.7 \times 10^{-3} = A e^{-5.225}$$

∴ 
$$A = 1.43 \text{ s}^{-1}$$

 $1.43 \, \mathrm{s}^{-1}$ 

Assign formal charges to all atoms of the thiocyanate ion, SCN-. Identify the most favored e. structure.

$$S = -1$$

$$N = -1$$

$$S = +1$$
  $N = -2$ 

$$S - C \equiv N$$

$$S = C = N$$

$$S - C \equiv N$$
  $S = C = N$   $S \equiv C - N$ 

#### 23. Long answer question (For full marks, you must show all work).

 $+ OH^- \rightleftharpoons H_2O + A^-$ 

8 marks

Meta-nitrobenzoic acid is a weak monoprotic acid with  $K_a = 6.5 \times 10^{-3}$ . Exactly fifty mL of 0.05 M meta-nitrobenzoic acid is titrated with 0.10 M NaOH. Calculate the pH:

a. Before any titrant is added.

b. After the addition of 10 mL of titrant

HA

[I] 2.5x10<sup>-3</sup>mol 1.0x10<sup>-3</sup>mol 0 
$$n_{HA} = 0.050Lx0.05mol/L = 2.5x10^{-3} mol$$
[C] -1.0x10<sup>-3</sup>mol -1.0x10<sup>-3</sup>mol +1.0x10<sup>-3</sup>mol  $n_{A-} = 0.010Lx0.010mol/L = 1.0x10^{-3} mol$ 
[Eq] 1.5x10<sup>-3</sup>mol 0 1.0x10<sup>-3</sup>mol 
$$pH = pK_a + log\left(\frac{[A^-]}{[HA]}\right) \qquad pH = 2.187 + log\left(\frac{[1.0x10^{-3}]}{[1.5x10^{-3}]}\right) \qquad pH = 2.187 - 0.176 = 2.011$$
Or 
$$K_a = \frac{[H_3O^+][A^-]}{[HA]} \qquad 6.5x10^{-3} = \frac{[H_3O^+][1.0x10^{-3}]}{[1.5x10^{-3}]} \qquad [H_3O^+] = 9.75x10^{-3} \qquad \text{Therefore, pH} = 2.01$$

Converting moles to moles/L (**M**) will give the same answer; [HA] = 0.025 M; [A<sup>-</sup>] = 0.01667 M  $n_{\rm HA} = 0.050 \, \rm Lx \, 0.050 \, Lx \, 0.050 \, Lx \, 0.050 \, L = 2.5 \, x \, 10^{-3} \, mol;$  [HA] = (2.5 x 10<sup>-3</sup> - 1.0 x 10<sup>-3</sup>) mol / 0.060 L = 0.025 M  $n_{\rm A-} = 0.010 \, \rm Lx \, 0.010 \, mol/L = 1.0 x \, 10^{-3} \, mol;$  [A<sup>-</sup>] = 1.0 x 10<sup>-3</sup> mol / 0.060 L = 0.01667 M

$$pH = 2.187 + log\left(\frac{0.01667}{0.0250}\right) = \ 2.011 \\ 6.5x10^{-3} = \frac{[H_3O^+][0.01667]}{[0.0250]} \ ; \ pH = 2.01$$

#### c. At the equivalence point

#### Volume of NaOH at equivalence point: $50 \times 0.05 / V? \times 0.010 = 25 \text{ mL}$

- the sodium salt of m-nitrobenzoic acid (Na A) will give the anion A, a conjugate base of the weak acid (nitrobenzoic acid).  $[A^-] = 50 \times 0.05 / 50 + 25 = 0.03333 \text{ M}$ 

$$A^{-}_{(aq)} + H_2O_{(l)} \rightleftharpoons OH^{-}_{(aq)} + HA_{(aq)}$$
  $K_b = K_w / K_a$ 
[Eq] 0.03333-x x x  $K_b = 1.0x10^{-14} / 6.5x10^{-3} = 1.54x10^{-12}$ 
 $K_b = 1.54x10^{-12} = \frac{(x)(x)}{0.0333 - x}$ ; x can be ignored!
 $x = [OH^{-}] = 2.2645x10^{-7}$  or  $pOH = 6.645$  and  $pH = 7.35$ 

#### d. After the addition of 40 mL of titrant

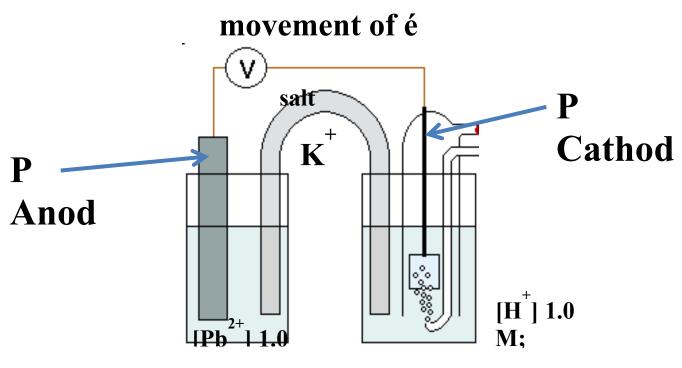
Total volume = 50 + 40 = 0.090 LExcess NaOH = 40.0 - 25.0 = 0.015 L[OH] =  $(0.015 L \times 0.10 \text{ mol/L}) / 0.090 L = 0.01666 M$ Therefore pOH = 1.78 and pH = 12.22

#### 24. Long answer question (For full marks, you must show all work).

The standard electrode potential for the  $Pb(s)/Pb^{2+}$  redox couple is -0.126 V. Sketch the cell in which this potential was measured.

#### **Identify:**

- a. All concentrations/pressures
- b. The cathode and the anode
- c. The direction in which cations move in the salt bridge
- d. The direction in which electrons would move if a light bulb were connected across the cell.



flow of cations to cathode movement of é anode to

For the species given below, draw Lewis (2D) and VSEPR (3D) structures and identify electron pair geometries, molecular geometries, bond angles around the central atom and molecular polarity. Enter your answers in the appropriate boxes:

6 Marks

Species	NCl <sub>3</sub>	XeOF <sub>4</sub>	SF <sub>4</sub>
Lewis structure (2D) (Show all valence electrons) Note: — = 2 é	;Ċi: - :Ċi:	.:	::::::::::::::::::::::::::::::::::::::
VSEPR structure (3D)	:Ċi	:E.W. Xe	F - S - F
Electro pair geometry	Tetrahedral	Octahedral	Trigonal bipyramidal
Molecular Geometry	Trigonal pyramidal	Square pyramidal	See saw
Bond angles	< 109.7°	< 90° & < 180°	<90°, <120° & <180°
Polar or Non-polar	Polar	Polar	Polar

#### Data Sheet – CHEM 209

<b>1</b> 1A							P	eriod	ic Tal	ble							<b>18</b> 8A
1 <b>H</b> 1.008	<b>2</b> 2A											13 3A	14 4A	15 5A	16 6A	17 7A	2 <b>He</b> 4.003
3	4											5	6	7	8	9	10
Li	Be											В	C	N	0	F	Ne
6.941	9.012											10.81	12.01	14.01	16.00	19.00	20.18
11	12											13	14	15	16	17	18
Na	Mg	3	4	5	6	7	8	9	10	11	12	Al	Si	P	$\mathbf{S}$	Cl	Ar
22.99	24.31											26.98	28.09	30.97	32.07	35.45	39.95
19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr
39.10	40.08	44.96	47.88	50.94	52.00	54.94	55.85	58.93	58.69	63.55	65.38	69.72	72.59	74.92	78.96	79.90	83.80
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe
85.47	87.62	88.91	91.22	92.91	95.94	(98)	101.1	102.9	106.4	107.9	112.4	114.8	118.7	121.8	127.6	126.9	131.3
55	56	57*	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Cs	Ba	La	Hf	Ta	$\mathbf{W}$	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn
132.9	137.3	138.9	178.5	180.9	183.9	186.2	190.2	192.2	195.1	197.0	200.6	204.4	207.2	209.0	(209)	(210)	(222)
87	88	89**	104	105	106	107	108	109	110	111							
Fr	Ra	Ac	Rf	Ha	Sg	Ns	Hs	Mt	Uun	Uuu							
(223)	226.0	(227)	(261)	(262)	(263)	(262)	(265)	(266)	(269)	(272)							

Lanthanides *	58	59	60	61	62	63	64	65	66	67	68	69	70	71
Lanthaniues	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu
	140.1	140.9	144.2	(145)	150.4	152.0	157.3	158.9	162.5	164.9	167.3	168.9	173.0	175.0
Actinides **	90	91	92	93	94	95	96	97	98	99	100	101	102	103
Actiniucs	Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lr
	232.0	231.0	238.0	237.0	(244)	(243)	(247)	(247)	(251)	(252)	(257)	(258)	(259)	(260)

Strong Acids: HCl, HBr, HI, HNO<sub>3</sub>, H<sub>2</sub>SO<sub>4</sub>, HClO<sub>4</sub>
Strong Bases: Hydroxides of Group 1A (Li to Cs) and Group 2A (Ca. Sr. B.

Strong Bases: Hydroxides of Group 1A (Li to Cs) and Group 2A	A (Ca, Sr, Ba)					
Constants:	Conversion factors:					
Gas constant, $R = 0.08205 \text{ L atm mol}^{-1} \text{ K}^{-1}$	$1 J = 1 kg m^2 s^{-2}$ $1 Pa = 1 kg m s^{-2}$					
$= 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$	$T K = T^{\circ}C + 273.15$					
Avogadro's number: $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$	1 L atm = 101.3 J					
Faraday: $F = 96,485 \text{ C} / \text{mol electrons}$	1atm = 760.0 torr = 101.3 kPa = 760.0 mm Hg = 1.013 bar					
Planck's constant $h = 6.626 \times 10^{-34} \text{Js}$	$1 L = 10^{-3} m^3$					
Speed of light, $c = 2.998x \ 10^8 \text{m/s}$	$1 \text{ C} = 1 \text{ J/V}$ $1 \text{ A} = 1 \text{ C s}^{-1}$					
Rydberg constant, $R = 1.09678 \times 10^{-7} \text{ m}^{-1}$	STP conditions: 0 °C, 1 atm					
Factoring constant, $R_H = Rhc = 2.18^{-18} J$	Electrochemical standard state: 1 atm, 1M, 25 °C					

SIP conditions: 0 °C, 1 atm Electrochemical standard state: 1 atm, 1M, 25 °C 
$$[AJ_t = -kt + [AJ_0] \qquad \ln[AJ_t = -kt + \ln[AJ_0] \qquad \frac{1}{[AJ_t]} = kt + \frac{1}{[AJ_0]} \qquad \ln\left(\frac{[AJ_0]}{[AJ_t]}\right) = kt$$

$$t_{1/2} = \frac{[AJ_0]}{2k} \qquad t_{1/2} = \frac{0.693}{k} \qquad t_{1/2} = \frac{1}{k[AJ_0]} \qquad k = Ae^{\frac{E_a}{RT}} \qquad \ln\left(\frac{K_2}{K_1}\right) = \frac{\Delta H}{R} \left(\frac{1}{T_1} - \frac{1}{T_2}\right)$$

$$\ln\left(\frac{k_2}{k_1}\right) = \frac{E_a}{R} \left(\frac{1}{T_1} - \frac{1}{T_2}\right) \qquad PV = nRT \qquad K_p = K_c(RT)^{An} \qquad ax^2 + bx + c = 0$$

$$pH = -\log[H^+] \qquad K_w = K_a K_b \qquad K_{sp} = 1/K_d \qquad K_f = 1/K_d \qquad x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$$

$$pH = pK_a + \log\left(\frac{[\text{cong. base}]}{[\text{cong. acid}]}\right) \qquad \text{or} \qquad pOH = pK_b + \log\left(\frac{[\text{cong. acid}]}{[\text{cong. base}]}\right)$$

$$E^\circ = E^\circ_{\text{cathode}} - E^\circ_{\text{anode}} \qquad E = E^\circ - \frac{0.0592}{n_e} \log Q \qquad E^\circ = \frac{0.0592}{n_e} \log K \qquad \text{or} \qquad nFE^\circ = RTlnK$$

$$q = It \qquad q = n_e F \qquad c = \lambda v \qquad E = hv \qquad E = mc^2 \qquad \frac{1}{\lambda} = R\left(\frac{1}{n_1^2} - \frac{1}{n_2^2}\right) \qquad \Delta E = -R_H\left(\frac{Z^2}{n_f^2} - \frac{Z^2}{n_i^2}\right)$$

$$E = -R_H\left(\frac{Z}{n}\right)^2 \qquad \text{or} \qquad E_n = -\frac{Rhc}{n^2} \qquad \text{for single electron species}$$