=======================================			=========	=========
12.1]. Which of th	e following liquids wo	uld make a good solv	ent for bromine, Br	2?
A. CCl ₄	B. H ₂ O	C. C	H₃OH	D. NH ₃
<mark>A]</mark>				
1.114 g/mL a	y mass solution of phosat 20°C. What is the mass of phosphoric acid	olarity of this solution		ity of
A. 0.0114 M	B. 0.0568 M	C. 0.114 M	D. 2.27 M	E. 5.51 M
A. 0.0114 M				
A. 0.0114 M				
D]		(C. H O.)		······································
D] 12.3]. What is the 1	molarity of 0.645 m gl f glucose is 180.2 g. Tl	0 12 0 1		vent: H ₂ O).
D] 12.3]. What is the 1	•	0 12 0 1		vent: H ₂ O). E. 0.737 M
D] 12.3]. What is the I The molar mass o A. 0.0251 M	f glucose is 180.2 g. Tl	he density of the solut	ion is 1.18 g/mL.	-
D] 12.3]. What is the I The molar mass o	f glucose is 180.2 g. Tl	he density of the solut	ion is 1.18 g/mL.	-
D] 12.3]. What is the I The molar mass o A. 0.0251 M	f glucose is 180.2 g. Tl	he density of the solut	ion is 1.18 g/mL.	<u>-</u> ·
D] [12.3]. What is the range of the molar mass of A. 0.0251 M [D]	f glucose is 180.2 g. Tl B. 0.1256 M	he density of the solut	ion is 1.18 g/mL. D. 0.682 M	E. 0.737 M
[12.4]. A mole fra is X= 0.2 . What	B. 0.1256 M action of solute (methat is the molality of this	c. 0.454 M C. 0.454 M anol: CH ₃ OH) in aque	D. 0.682 M cous solution (Solver	E. 0.737 M
[12.4]. A mole fra is X= 0.2 . What	f glucose is 180.2 g. Tl B. 0.1256 M action of solute (metha	c. 0.454 M C. 0.454 M anol: CH ₃ OH) in aque	D. 0.682 M cous solution (Solver	E. 0.737 M

[12.5	[5]. Which respond	onse lists a	ll the follow	wing pairs tha	t are mis	cible liquids.	
		Pair #2:	acetic aci	C8H18) and wated (CH3COOH C8H18) and car) and wa		4)
<u>[E]</u>	A) 1	, 3 B)	1, 2	C) 3	D) 2	E) 2,	3
[12.6	[]. An exothermic	solution p	rocess is	described by	which of	the following	?
[D]		B) ΔH > 0		feels hot		0) ΔH < 0, so E) ΔH = 0	lution feels hot
[12.7				n a solution m			g of KI in 5.8 g of
[A]	A) 0.060	0 B	0.064	C) 0.37	D) 0.59	E) 6.4
[12.8		mass soluti	on of acet	` '			of 0.9849 g/mL at
	A) 0.621 M	B) 1.	.61 M	C) 1.66 I	М	D) 1.71 M	E) 16.9 M
[B]							
	9]. A solution th mole fraction of the		6 by mass	of phenol (Ce	 вН5ОН) ir	n water is pre	epared. What is the
	A) 0.0969 E	3) 0.0643	C) 0.08	83 D) 0.06	604	E) 0.357	
[C]							
[12.	10]. What is the KBr, X, equal	•	an aqueo	ous KBr solution	on with a	mole fraction	n of
[D]	A) 0.325 <i>m</i>	B) 0.68	32 m	C) 13.6 m	D) 18	3.0 <i>m</i> E	i) 29.2 m

in v	vater at 25°C if the	-	essure is 741 mm		the solubility of O ₂ (g) law constant
	A) 2.7 × 1 B) 1.3 × 1 C) 6.2 × 1	0^{-3} M	D) 9.6 × 7 E) 0.96 N		
[A]					
[12.12			at 20°C is 17.5 mi 00 × 10 ² g of sucr		vapor pressure of water and 3.50 × 10 ² g
[C]	A) 0.51 m B) 16.0 m C) 17.0 m	nmHg	D) 18.0 mmHg E) 19.4 mmHg		
[12.13		oiling point of a so 2O) = 0.52 °C/m)	lution of 11.0 g of	lactose (C12H22O	 11) in 145.0g of
[C]	A) 0.1°C	B) 99.9 °C	C) 100.1°C	D) 102.2 °C	E) 105.7 °C
	of a solution		:(СзНтОН, 60.10 g/ ressure of 25 atm :		eeded to make 750 mL 0821 L·atm/K·mol)
[46 g]					
		nms of water are re a of a 0.452 m solution		27.8 g of ammon	nium nitrate NH4NO3
	A) 769 g	B) 36.2 g	C) 100. g	D) 0.157 g	E) 157 g
Ans: A	·				

[12.16] A solution is 40.0% by mass be vapor pressure of pure benzene and pummHg and 91.32 mmHg, respectively. 20°C:	ure carbon tétrachlo	oride at this temp	peraturé is 74.61
A) 66.4 mmHg B) 73.8 mmHg C) 81.8 mmHg		D) 83.0 mmHg E) 84.6 mmHg	
[Ans: C]			
[12.17] A solution of chloroform, CHC from Raoult's law. This result im	•	H ₃) ₂ CO, exhibits	a negative deviation
A) chloroform-chloroform interactions a	re stronger than ch	loroform-aceton	ie
interactions. B) chloroform-chloroform interactions a	ire weaker than chlo	oroform-acetone	<u> </u>
interactions.			
C) acetone-acetone interactions are strD) acetone-acetone interactions are weE) Both B and D.F) Both A and C.G) Both A and D.			
[Ans: E]			
[12.18] Which of the following aqueous dissociation for all soluble ionic		highest boiling p	point (assume 100%
A) 0.10 <i>m</i> AI(NO ₃) ₃ B) 0.11 <i>m</i> Na ₂ SO ₄ C) 0.15 <i>m</i> K ₂ CO ₃	D) 0.18 <i>m</i> NaCl E) 0.35 <i>m</i> C ₆ H ₁₂	O ₆	
[Ans: C]			
[12.19] When 24.0 g of glucose (a non has a freezing point of –0.47°C. What i 1.86°C/m.	• ,	•	
A) 41.9 g B) 47.5 g	C) 54.9 g	D) 178 g	E) 190. g
[Ans: E]			

solution consist	nzene, C6H6, freeze ing of cyclohexane ().0°C? (For benzene Kb = 2.79°C/m)	C ₆ H ₁₂) dissolved	in benzene if the	freezing point of	nt of a
A) 77.4°C	B) 80.9°C	C) 82.8°C	D) 83.1°C	E) 91.2°C	
[Ans: C]					
pressure of 17.0	eous fructose solution at 25°C. Find the Kf = 1.86°C/m; mo	he temperature a	at which this solut	ion freezes.	n osmotic
A) -1.30°C	B) -1.41°C	C) -1.52	2°C D) –	1.57°C E) –	1.69°C
[Ans: B]					
	oor pressure of water 1.50 <i>m</i> K ₂ CO ₃ at this				ssure of a
	A) 1.89 mmHg B) 5.39 mmHg C) 66.5 mmHg	g E) 108	mmHg mmHg		
[Ans: C]					
isolated from benzene,	ne, an important how om the thyroid gland the freezing point of K _f of 5.12°C/m. Wha	. If 0.455 g of thy the solution is 5.	roxine is dissolve 144°C. Pure benz	d in 10.0 g of	•
[Ans: 777 g/mol]				
	following solutions i Sc(NO ₃) ₃ , 0.15 <i>m</i> K		 sing melting poin	t: 0.20 <i>m</i> glycerol,	
[Ans: glycerol <	Sc(NO ₃) ₃ < K ₂ CO ₃]				

[12.25]. Which of the CH ₃ COONa (moderate The density of the control of the cont	olar mass = 82	2.0 g/mol) in		% by mass sol	ution of sodium ace	etate,
A) 2.26	$6 \times 10^{-6} \text{ M}$	3) 0.207 M	C) 2.07 M	D) 2.26 M	E) 2.72 M	
Ans: D						
[12.26]. What is th of Na ₂ SO ₄ , X =	•	mass of a N	a ₂ SO ₄ solution	n in water with a	a mole fraction	
A) 2.76%	B) 4.2	25%	C) 19.1%	D) 73.4 %	E) 80.9%	
Ans: E						
[12.27] The vapor p water over a solution this temperature?						of
A) 1.22 m B) 42.3 m C) 43.0 m	ımHg		D) 70.7 mmHg E) 73.2 mmHg			
Ans: D						
[12.28] . What is the of H ₂ O?	ne freezing po Krfor water is		tion that conta	ains 10.0 g of gl	lucose (C6H12O6) ir	n 100.g
A) +0.10°C	B) +0.	186°C	C) -0.10°C	D) -0.186	6°C E) –1.03	3°C
Ans: E						
[12.29]. How many a solution with a						 .5 L of
A) 0.57 g	B) 6.8 g	C) 2.0 ×	10^2 g D	$) 4.9 \times 10^{2} g$	E) 5.8×10^3	g
Ans: D						

	 A) 727 g/mol B) 663 g/mol C) 1.10 x 10² g/mol 	D) 1.15 g/mol E) 0.872 g/mol
Ans: B		
	Assuming 100% dissocn its van't Hoff factor i?	iation, which of the following compounds is listed incorrectly
	A) Na ₂ SO ₄ , <i>i</i> = B) NH ₄ NO ₃ , <i>i</i> = C) Sucrose, <i>i</i> =	E) Mg(NO ₃) ₂ , $i = 3$
Ans: D		
[12.32].	During osmosis	
	B) pure solutes diffuse th C) pure solvent and a so	through a membrane but solutes do not. Irough a membrane but solvent does not. Iution both diffuse at the same time through a membrane. a membrane into a solution and build up pressure.

Chapter 13: Chemical Kinetics

[13.1]. Chlorine dioxide reacts in basic water to form chlorite and chlorate according to the following chemical equation:

$$2CIO_2(aq) + 2OH^-(aq) \rightarrow CIO_2^-(aq) + CIO_3^-(aq) + H_2O(I)$$

Under a certain set of conditions, the initial rate of disappearance of chlorine dioxide was determined to be 2.30×10^{-1} M/s. What is the initial rate of appearance of chlorite ion (ClO₂) under those same conditions?

A)
$$5.75 \times 10^{-2} \text{ M/s}$$

$$\dot{\text{C}}$$
) 2.30 x 10⁻¹ M/s

[B]

[13.2]. The reaction $A + 2B \rightarrow \text{products}$ has been found to have the rate law, rate = k[A] [B] 2 . While holding the concentration of A constant, the concentration of B is increased from x to 3x. Predict by what factor the rate of reaction increases.

A) 3

B) 6

C) 9

D) 27

E) 30

[C]

[13.4]. The reaction A + 2B \rightarrow products was found to follow the rate law: rate = k [A] 2 [B]. Predict by what factor the rate of reaction will increase when the concentration of A is doubled, the concentration of B is tripled, and the temperature remains constant.

A) 5

B) 6

C) 12

D) 18

E) None of these.

[C]

[13.5] . It takes 42.0 min for the concentration of a reactant in a first-order reaction to drop from 0.45 M to 0.32 M at 25°C. How long will it take for the reaction to be 90% complete?

A) 13.0 min

B) 86.0 min

C) 137 min

D) 222 min

E) 284 min

[E]

[13.6]. Use the following data to determine the rate law for the reaction shown below.

$$2NO + H_2 \rightarrow N_2O + H_2O$$

Expt. #	[NO] ₀	[H ₂] ₀	Initial rate	
1	0.021	0.065	1.46 M/min	
2	0.021	0.260	1.46 M/min	
3	0.042	0.065	5.84 M/min	

- rate = k [NO] D) rate = k [NO] 2 [H₂] rate = k [NO] 2 E) rate = k [NO] 4 [H₂] 2

rate = k [NO] [H₂]

[B]

[13.7]. The isomerization of cyclopropane to form propene is a first-order reaction.

At 760 K, 85% of a sample of cyclopropane changes to propene in 79.0 min. Determine the rate constant for this reaction at 760 K.

C) 2.42

[E]

- [13.8]. A city's water supply is contaminated with a toxin at a concentration of 0.63 mg/L. Fortunately, this toxin decomposes to a safe mixture of products by first-order kinetics with a rate constant of 0.27 day⁻¹. How long will it take for half of the toxin to decompose?
- [E]
- A) 0.17 days B) 0.27 days C) 0.38 days D) 2.3 days
- E) 2.6 days

[13.9]. The isomerization of cyclopropane to form propene is a first-order reaction. At 760 K, 15% of a sample of cyclopropane changes to propene in 6.8 min. What is the half-life (the time required for the half of the initial amount of the reactant to be converted into the product) of cyclopropane at 760 K?

- A) 3.4×10^{-2} min
- B) 2.5 min
- C) 23 min
- D) 29 min
- E) 230 min

[D]

- The activation energy for the reaction CH₃CO → CH₃ + CO is 71 kJ/mol. How many [13.10]. times greater is the rate constant for this reaction at 170°C than at 150°C?
 - A) 0.40
- B) 1.1
- C) 2.5
- D) 4.0
- E) 5.0

[C]

The isomerization of methyl isocyanide, $CH_3NC \rightarrow CH_3CN$, follows first-order kinetics. [13.11]. The half-lives were found to be 161 min at 199°C and 12.5 min at 230°C.

Calculate the activation energy for this reaction.

A) 6.17×10^{-3} kJ/mol

D) 124 kJ/mol

B) 31.4 kJ/mol

C) 78.2

E) 163 kJ/mol

[E]

The rate law for the reaction $2NO_2 + O_3 \rightarrow N_2O_5 + O_2$ is rate = $k[NO_2][O_3]$. [13.12] Which one of the following mechanisms is consistent with this rate law?

A)
$$NO_2 + NO_2 \rightarrow N_2O_4$$
 (fast)
 $N_2O_4 + O_3 \rightarrow N_2O_5 + O_2$ (slow)

B) NO₂ + O₃
$$\rightarrow$$
 NO₅ (fast)
NO₅ + NO₅ \rightarrow N₂O₅ + 5/2O₂ (slow)

C)
$$NO_2 + O_3 \rightarrow NO_3 + O_2$$
 (slow)
 $NO_3 + NO_2 \rightarrow N_2O_5$ (fast)

D)
$$NO_2 + NO_2 \rightarrow N_2O_2 + O_2$$
 (slow)
 $N_2O_2 + O_3 \rightarrow N_2O_5$ (fast)

Ans: Cl

[13.13]	catalyst,	the Eais	55 kJ/mol. F	How many time	s faster is th	ne catalyzed	In the presence of a than the mains the same.
[Ans: A		5.0 times	B) 1.16 t	imes C) 15	times D)	2.0 times	E) 0.2 times
[13.14]	 What is t	the rate la	aw that corre	sponds to the	data shown	for the reacti	 on 2A + B → C?
		Exp.		Initial [B]		te	
		1 2 3 4	0.015 0.030 0.060 0.060	0.022 0.044 0.044 0.066			
Rate =	k[B] ² .						
[13.15]	For the a straig		whose rate	law is rate = k[X], a plot of	which of the	following is
[B]	A) B) C)	ln [X] v	ersus time ersus time ersus time		D) [X] versus E) In [X] vers		
			-	und to follow the 1.55 M, how i			where for [A]t = 0.150M
	,	A) 6.58 s	B) 395 s	C) 6.02 s	D) 17.0 s	E) 1.02 x 1	0^3 s
[E] 							

[13.17] The thermal decomposition of acetaldehyde, CH₃CHO \rightarrow CH₄ + CO, is a second-order reaction. The following data were obtained at 518°C.

time, s	Pressure CH ₃ CHO, mmHg
0	364
42	330
105	290
720	132
42	330
105	290

Based on the data given, what is the half-life for the disappearance of acetaldehyde?

- A) 1.5×10^5 s
- B) 410 s C) 5.4×10^7 s
- D) 520 s
- E) 305 s



- [13.18] Concerning the rate law, Rate = $k[A]_0$, what are appropriate units for the rate constant k?
- A) s^{-1} B) $M^{-1}s^{-1}$ C) $M^{-2}s^{-1}$ D) M/s E) M^2/s



- [13.19] At 25°C, the second-order reaction NOCl(g) \rightarrow NO(g) + 1/2Cl₂(g) is 50% complete after 5.82 hours when the initial concentration of NOCI is 4.46 mol/L. How long will it take for the reaction to be 75% complete?
 - A) 8.22 hr
- B) 11.6 hr C) 15.5 hr D) 17.5 hr E) 23.0 hr

[D]

[13.20]

The first-order rate constant for the decomposition of dimethyl ether

$$H_3C$$
 CH_3 $CH_4(g) + H_2(g) + CO(g)$,

is 3.2×10^{-4} s⁻¹ at 450 °C. The reaction is carried in a constant volume flask. Initially only dimethyl ether is present and the pressure is 0.350 atm. What is the pressure of the system after 8 min. Assume ideal behavior.

[0.45 atm]

[13.21] For the chemical reaction A \rightarrow C, a plot of 1/[A] _t versus time was found to give a straight line with a positive slope. What is the order of reaction?								
B) 1 C) :	A) zeroth B) first C) second D) Such a plot cannot reveal the order of the reaction.							
[C]								
by wh	[13.22]. The reaction A + 2B → products was found to have the rate law, rate = k[A] [B] ² . Predict by what factor the rate of reaction will increase when the concentration of A is doubled and the concentration of B is also doubled.							
	A) 2	B) 4	C) 6	D) 8	E) 9			
Ans: D								
[13.23]. Concerning the rate law, Rate = k[A][B], what are appropriate units for the rate constant k?								
	A) s ⁻¹	B) M ⁻¹ s ⁻¹	C) M ⁻² s ⁻¹	D) M/s		E) M^2/s		
Ans: B								

Chlorine dioxide reacts in basic water to form chlorite and chlorate according to the [13.24]. following chemical equation:

$$2CIO_2(aq) + 2OH^-(aq) \rightarrow CIO^{2-}(aq) + CIO^{3-}(aq) + H_2O(I)$$

A kinetic study of this reaction under a certain set of conditions yielded the data below.

Ехр	[CIO ₂] (M)	[OH ⁻] (M)	-Δ[ClO ₂] /Δt (M/s)
1	0.0500	0.100	5.75 x 10 ⁻²
2	0.100	0.100	2.30 x 10 ⁻¹
3	0.100	0.0500	1.15 x 10 ⁻¹

Which one of the following is the rate law for this reaction?

- A) rate = $k[ClO_2] [OH^{-}]$ B) rate = $k[ClO_2]^{2} [OH^{-}]$
- D) rate = $k[ClO_2]^2[OH^2]^2$
- E) rate = $k[ClO_2]^4[OH^-]$
- C) rate = $k[ClO_2] [OH^-]^2$

Ans: B

[13.25]. At 25°C the rate constant for the first-order decomposition of a pesticide solution is 6.40×10^{-3} min⁻¹. If the starting concentration of pesticide is 0.0314 M, what concentration will remain after 62.0 min at 25°C?

- A) 1.14×10^{-1} M D) 2.11×10^{-2} M E) 2.68×10^{-2} M

B) 47.4 M

E) $2.68 \times 10^{-2} \text{ M}$

C) -8.72.0 M

Ans: D

[13.25]. The Arrhenius equation is $k = Ae^{-Ea/RT}$. The slope of a plot of ln k vs. 1/T is equal to

- A) -k B) k C) E_a D) $-E_a/R$
- E) A

Ans: D

[13.26]. The activation energy for the following first-order reaction is 102 kJ/mol.

$$N_2O_5(g) \rightarrow 2NO_2(g) + 1/2O_2(g)$$

The value of the rate constant (k) is $1.35 \times 10^{-4} \,\mathrm{s}^{-1}$ at 35° C. What is the value of k at 0°C?

A)
$$8.2 \times 10^{-7} \text{ s}^{-1}$$

D)
$$2.2 \times 10^{-2} \,\mathrm{s}^{-1}$$

B)
$$1.9 \times 10^{-5} \text{ s}^{-1}$$

$$\dot{\text{C}}$$
 4.2 × 10⁻⁵ s⁻¹

Ans: A

[13.27]. The following mechanism has been suggested for the reaction:

$$H_2O_2 + 2H^+ + 2I^- \rightarrow I_2 + 2H_2O$$

$$H_2O_2 + I \longrightarrow HOI + OH^-$$
 Slow
 $OH^- + H^+ \longrightarrow H_2O$ Fast
 $HOI + H^+ + I^- \longrightarrow I_2 + H_2O$ Fast

Identify the rate law that is consistent with this mechanism

- A) Rate = $k[H_2O_2][I^{-}]$
- D) Rate = $k[HOI][H^+][I^-]$
- B) Rate = $k[HOI][OH^{-}]$ E) Rate = $k[H_2O_2][I^{-}]^2[H^{+}]^2$
- C) Rate = $k[OH^{-}][H^{+}]$

Ans: A Category: Medium

[13.28]. The reaction $2A \rightarrow products$ is second order with respect to A. If the concentration of A drops from 1.05 M to 0.815 M in a time of 15.0 min, what is the rate constant for this reaction (the same time units may be used)?

Ans: 1.83 x 10-2 M-1min-1

[13.29]. For what order reaction does the half-life get longer as the initial concentration increases?

- A) zero order
- B) first order
- C) second order
- D) none of them because half-life is always independent of the initial concentration

Ans: A

[13.30]. At 30°C, by how much is a reaction's activation energy decreased by the addition of a catalyst if the catalyst triples the reaction rate?

- A) 2.77 kJ/mol B) 274 J/mol C) 2.70 J/mol

- D) 119 J/mol
- E) 1.20 kJ/mol

Ans: A

[13.31]. At 700 K, the rate constant for the following reaction is $6.2 \times 10^{-4} \, \text{min}^{-1}$.

How many minutes are required for 20% of a sample of cyclopropane to isomerize to propene?

- A) 1,120 min B) 360 min C) 3710 min D) 1.4×10^{-4} min E) 280 min

Ans: B

[13.32] The isomerization of cyclopropane follows first order kinetics. The rate constant at 700 K is $6.20 \times 10^{-4} \text{ min}^{-1}$, and the half-life at 760 K is 29.0 min.

Calculate the activation energy for this reaction.

- A) 5.07 kJ/mol
- D) 60. kJ/mol
- B) 27.0 kJ/mol
- E) 270. kJ/mol
- C) 50.7 kJ/mol

Ans: E

Chapter 14: Chemical Equilibrium

[14.1] . A reaction with an equilibrium constant $K_c = 1.5 \times 10^{21}$ would consist of which of the following at equilibrium:

- A) approximately equal reactants and products
- B) some reactants and products with reactants slightly favored
- C) some reactants and products with products slightly favored
- D) essentially all reactants
- E) essentially all products

Ans: E

[14.2]. The solubility of silver chloride can be increased by dissolving it in a solution containing ammonia.

AgCl (s)
$$\rightarrow$$
 Ag⁺ (aq) + Cl⁻ (aq) K₁ = 1.6 x 10⁻¹⁰
Ag⁺ (aq) + 2NH₃ (aq) \rightarrow Ag(NH₃)₂⁺ (aq) K₂ = 1.5 x 10⁷

What is the value of the equilibrium constant for the overall reaction?

AgCl (s) + 2NH₃ (aq)
$$\stackrel{\blacktriangleright}{\longleftarrow}$$
 Ag(NH₃)₂⁺ (aq) + Cl⁻ (aq) K_{net} = ?
A) 1.5 x 10⁷ B) 2.4 x 10⁻³ C) 3.1 x 10⁻³ D) 2.3 x 10¹⁴ E) 2.4 x 10⁷

Ans: B

[14.3]. 2.50 mol NOCl was placed in a 2.50 L reaction vessel at 400°C. After equilibrium was established, it was found that 28% of the NOCI had dissociated according to the equation 2NOCI(q) $2NO(g) + Cl_2(g)$. Calculate the equilibrium constant, Kc, for the reaction. A) 0.021 B) 0.039 C) 0.169 D) 26 E) 47 Ans: A [14.4]. Equilibrium is established for the reaction $2X(s) + Y(g) \rightleftharpoons 2Z(g)$ at 500K, $K_c = 100$. Determine the concentration of Z in equilibrium with 0.2 mol X and 0.50 M Y at 500K. A) 3.2 M B) 3.5 M C) 4.5 M D) 7.1 M E) None of these. Ans: D [14.5]. At 400°C, $K_c = 64$ for the equilibrium $H_2(g) + I_2(g) \stackrel{\longrightarrow}{\longleftarrow} 2HI(g)$. If 3.00 mol H_2 and 3.00 mol l2 are introduced into an empty 4.0 L vessel, find the equilibrium concentration of HI at 400°C. A) 0.15 M B) 1.2 M C) 2.4 M D) 4.8 M E) 5.8 M

Ans: B

[14.6] For the following reaction at equilibrium in a reaction vessel, which change will cause

[14.6]. For the following reaction at equilibrium in a reaction vessel, which change will cause the Br₂ concentration to *decrease*?

$$2NOBr(g) \stackrel{\longrightarrow}{\longleftarrow} 2NO(g) + Br_2(g), \Delta H^{o}_{rxn} = 30 \text{ kJ/mol}$$

- A) Increase the temperature.
- B) Remove some NO.
- C) Add more NOBr.
- D) Compress the gas mixture into a smaller volume.
- E) Add a catalyst

Ans: D

[14.7]. Consider the following equilibrium,

$$4NH_3(g) + 3O_2(g)$$
 \longrightarrow $2N_2(g) + 6H_2O(g) + 1531 kJ$

State whether the concentrations of the reactants would increase, decrease, or remain constant when the temperature is increased.

Ans: increase

[14.8]. 4.21 moles of S₂Cl₄ are introduced into a 2.0 L vessel.

$$S_2Cl_4(g)$$
 \longrightarrow $2SCl_2(g)$

At equilibrium, 1.25 moles of S₂Cl₄ are found to remain in the container. Calculate K_c for this reaction.

Ans: $K_c = 14.0$

[14.9]. The equilibrium constant expression for the reaction $2BrF_5(g) \rightarrow Br_2(g) + 5F_2(g)$ is

A)
$$K_c = [Br_2] [F_2] / [BrF_5]$$

D)
$$K_c = [BrF_5]^2 / [Br_2][F_2]^5$$

B)
$$K_c = [Br_2] [F_2]^5 / [BrF_5]^2$$

A)
$$K_c = [Br_2] [F_2] / [BrF_5]$$

B) $K_c = [Br_2] [F_2]^5 / [BrF_5]^2$

C) $K_c = [Br_2] [F_2]^2 / [BrF_5]^5$

D) $K_c = [Br_5]^2 / [Br_2] [F_2]^5$

E) $K_c = 2[BrF_5]^2 / ([Br_2] \times 5[F_2]^5)$

C)
$$K_c = [Br_2] [F_2]^2 / [BrF_5]^2$$

Ans: B

[14.10] Carbon tetrachloride reacts at high temperatures with oxygen to produce two toxic gases, phosgene and chlorine.

$$CCl_4(g) + 1/2O_2(g) \leftarrow COCl_2(g) + Cl_2(g)$$
, $K_c = 4.4 \times 10^9$ at 1,000 K

Calculate K_c for the reaction $2CCl_4(g) + O_2(g) \rightleftharpoons 2COCl_2(g) + 2Cl_2(g)$.

A)
$$4.4 \times 10^9$$

B)
$$8.8 \times 10^{9}$$

A)
$$4.4 \times 10^9$$
 B) 8.8×10^9 C) 1.9×10^{10} D) 1.9×10^{19} E) 2.3×10^{-10}

D)
$$1.9 \times 10^{19}$$

E)
$$2.3 \times 10^{-10}$$

Ans: D

I14.111 1.25 moles of NOCI were placed in a 2.50 L reaction chamber at 427°C. After equilibrium was reached, 1.10 moles of NOCI remained. Calculate the equilibrium constant, Kc, for the reaction

2NOCl(g) \longrightarrow $2NO(g) + <math>Cl_2(g)$.

A) 3.0×10^{-4} B) 1.8×10^{3} C) 1.4×10^{-3} D) 5.6×10^{-4} E) 4.1×10^{-3}

Ans: D

[14.12] At 250°C, the equilibrium constant K_p for the reaction $PCl_5(g) \nearrow PCl_3(g) + Cl_2(g)$ is 1.80. Sufficient PCIs is put into a reaction vessel to give an initial pressure of 2.74 atm at 250°C. Calculate the pressure of PCI₅ after the system has reached equilibrium.

A) 1.50 atm

- B) 1.24 atm C) 4.24 atm
- D) 0.94 atm
- E) 1.12 atm

Ans: B

- [14.13] For the reaction at equilibrium $2SO_3 \rightleftharpoons 2SO_2 + O_2 (\Delta H^0_{rxn} = 198 \text{ kJ/mol}),$ increasing the reaction temperature would:
 - A) Shift the equilibrium to the right and increase the value of the equilibrium constant K
 - B) Shift the equilibrium to the left and increase the value of the equilibrium constant K
 - C) Shift the equilibrium to the right and decrease the value of the equilibrium constant K
 - D) Shift the equilibrium to the left and decrease the value of the equilibrium constant K
 - E) No change

Ans: A

[14.14] 4.21 moles of S₂Cl₄ are introduced into a 2.0 L vessel.

$$S_2Cl_4(g) \implies 2SCl_2(g)$$

At equilibrium, 1.25 moles of S₂Cl₄ are found to remain in the container. Calculate K_c for this reaction.

Ans: $K_c = 14.0$

[14.15] 75.0 g of PCl₅(g) is introduced into an evacuated 3.00 L vessel and allowed to reach equilibrium at 250°C.

$$PCI_5(g) \longrightarrow PCI_3(g) + CI_2(g)$$

If $K_p = 1.80$ for this reaction, what is the total pressure inside the vessel at equilibrium?

- A) 2.88 atm
- B) 2.27 atm
- C) 4.54 atm
- D) 7.42 atm
- E) 9.69 atm

Ans: D

Chapter 15: Acids and Bases

[15.1]. In the reaction: $CH_3COOH(aq) + NH_2^-(aq) \leftarrow CH_3COO^-(aq) + NH_3(aq)$, the conjugate acid-base pairs are:

A) pair 1: CH₃COOH and CH₃COO⁻; pair 2: NH₂⁻ and NH₃ pair 2: NH₂ and CH₃COC pair 2: NH₃ and CH₃OO B) pair 1: CH₃COOH and NH₃; NH₂ and CH₃COO C) pair 1: CH₃COOH and NH₂⁻; D) pair 1: CH₃COOH and CH₃COO⁻; pair 2: NH₄⁺ and NH₃

NH₂ and NH₃ + E) pair 1: CH₃COOH and CH₃COO⁻; pair 2:

Ans: A

[15.2]. Identify the conjugate acid of SO₄²⁻

A) H₂SO₄

B) HSO₄

C) H₂SO₃

D) H₃O⁺ E) SO₃ ²⁻

Ans: B

[15.3]. The OH⁻ concentration in a 2.5×10^{-3} M Ba(OH)₂ solution is

A) 4.0×10^{-12} M.

D) 1.2×10^{-2} M.

B) 2.5×10^{-3} M.

E) 0.025 M.

C) 5.0×10^{-3} M.

Ans: C

[15.4] Which of the following solutions is acidic?

A) $[OH^{-}] = 1.0 \times 10^{-7} M$

D) $[H_3O^+] = 1.0 \times 10^{-10} M$

B) $[OH^{-}] > 1.0 \times 10^{-7} M$

E) $[H_3O^+] < 1.0 \times 10^{-7} M$

C) $[OH^{-}] = 1.0 \times 10^{-10} M$

Ans: C

[15 . 5]	Calculate the pH of a 0.14 M HNO ₂ solution that is 5.7% ionized.								
	A) 0.85	B) 1.70	C) 2.	10 D)	11.90	E) 13.10			
Ans: (
[15.6]	Calculate the h	ydrogen ion c	oncentration	in a solution ha	ving a pH of	4.60.			
A) 4.0×10^{-3} M D) 2.5×10^{-5} M B) 4.0×10^{-9} M E) 2.5×10^{-4} M C) 4.0×10^{-10} M									
Ans: I	O								
[15.7]	A 5.2 L sample solution. What is			s mixed with 2.3	3 L of a 0.20	M Sc(OH)з			
	A) 13.67	B) 13.89	C) 14.11	D) 14.23	E) 13.9	98			
Ans: I	=								
[15.8]	Arrange the ac	cids HBr, H ₂ Se	e, and H₃As ir	n order of increa	asing acid str	ength.			
	· · · · · · · · · · · · · · · · · · ·				H ₂ Se < HBr HBr < H ₂ Se				
Ans: [O.								
 [15.9]	Which one of these net ionic equations represents the reaction of a strong acid with weak base?								
A) $H^{+}(aq) + OH^{-}(aq) \rightarrow H_{2}O(aq)$ B) $H^{+}(aq) + CH_{3}NH_{2}(aq) \rightarrow CH_{3}NH_{3}^{+}(aq)$ C) $OH^{-}(aq) + HCN(aq) \rightarrow H_{2}O(aq) + CN^{-}(aq)$ D) $HCN(aq) + CH_{3}NH_{2}(aq) \rightarrow CH_{3}NH_{3}^{+}(aq)$						CN [−] (aq)			
Ans: [3								

[15.10]. The pH of a 0.14 M solution of an unknown monoprotic acid is 5.85. Calculate the K_a of the acid.

- A) 1.4×10^{-6} B) 1.4×10^{-11} C) 1.0×10^{-5} D) 7.1×10^{-9} E) 2.0×10^{-7}

Ans: B

[15.11]. The hydrolysis of NH₄F will result in which of the following types of solutions given:

$$(K_a(NH_4^+) = 5.6 \times 10^{-10}, K_b(F^-) = 1.4 \times 10^{-11})$$

$$K_b(F^-) = 1.4 \times 10^{-11}$$

- A) acidic, pH < 7 B) basic, pH > 7

C) neutral, pH = 7

Ans: A

[15.12]. The equilibrium constant for the reaction

 $C_6H_5COOH(aq) + CH_3COO^-(aq) \leftarrow C_6H_5COO^-(aq) + CH_3COOH(aq)$

is 3.6 at 25°C. If K_a for CH₃COOH is 1.8 × 10^{-5} , what is the acid dissociation constant for C₆H₅COOH?

- A) 5.0×10^{-6} B) 6.5×10^{-5} C) 2.3×10^{-4} D) 8.3×10^{-5} E) 5.6×10^{-6}

Ans: B

[15.13]. What mass of sodium formate (HCOONa) must be added to 350. mL of water in order to obtain a solution having a pH of 8.50? $[K_a(HCOOH) = 1.77 \times 10^{-4}]$

- A) 0.23 g B) 4.3 g
- C) 35 g
- D) 12 g
- E) 130 g

Ans: B

[15.14]. Write the formula for the conjugate acid of HPO₄ ²⁻

Ans: H₂PO₄

[15.15] The oxides CO₂ and SO₃ will form the following acids in water, respectively.

- A) H₂CO₃(aq) and H₂SO₃(aq)
- D) $H_2CO_2(aq)$ and $H_2SO_4(aq)$
- B) $H_2CO_3(aq)$ and $H_2SO_4(aq)$
- E) HCOO (aq) and H2SO3(aq)
- C) $H_2CO_2(aq)$ and $H_2SO_3(aq)$

Ans: B

[15.16] Which of the following is both a Lewis Acid and Brønsted Acid?

- A) BF₃
- B) NH₃ C) CO₂
- D) HNO₂
- E) PCI3

[15.17]. Consider the two gaseous equilibria:

$$SO_2(g) + 1/2O_2(g)$$
 SO₃(g)

 K_2

$$2SO_3(g) \implies SO_2(g) + O_2(g)$$

The values of the equilibrium constants K₁ and K₂ are related by

A)
$$K_2 = K_1^2$$

B)
$$K_2^2 = K_1^2$$

C)
$$K_2 = 1/K_1^2$$

D)
$$K_2 = 1/K_1$$

A) $K_2 = K_1^2$ B) $K_2^2 = K_1$ C) $K_2 = 1/K_1^2$ D) $K_2 = 1/K_1$ E) none of these.

Ans: C

[15.18]. When the following reaction is at equilibrium, which of these relationships is always true?

$$2NOCI(g) \rightleftharpoons 2NO(g) + CI2(g)$$

A) [NO] $[Cl_2]$ = [NOCI]

D) $2[NO] = [Cl_2]$

B) $[NO]_2[Cl_2] = [NOCl]^2$

E) $[NO]^2$ $[Cl_2] = K_c [NOCl]^2$

C) [NOCI] = [NO]

Ans: E

[15.19].

The brown gas NO₂ and the colorless gas N₂O₄ exist in equilibrium, 2NO₂ ADO₄. In an experiment, 0.625 mole of N2O4 was introduced into a 5.00 L vessel and was allowed to decompose until equilibrium was reached. The concentration of N2O4 at equilibrium was 0.0750 M. Calculate Kc for the reaction.

A) 7.5

B) 0.125

C) 0.0750

D) 0.10

E) 0.050

Ans: A

[15.20].

Consider the reaction $N_2(g) + O_2(g)$ \rightleftharpoons 2NO(g), for which $K_c = 0.10$ at 2,000°C. Starting with initial concentrations of 0.040 M of N2 and 0.040 M of O2, determine the equilibrium concentration of NO.

A) 5.4×10^{-3} M

B) 0.0096 M C) 0.013 M D) 0.080 M

E) 0.10 M

Ans: C

[15.21] Concerning the following reaction at equilibrium:

 $3Fe(s) + 4H_2O(g)$ Fe₃O₄(s) + $4H_2(g)$, increasing the concentration of the Fe(s) would:

- A) Shift the equilibrium to the right
- B) Shift the equilibrium to the left
- C) Increase the value of the equilibrium constant, K
- D) Decrease the value of the equilibrium constant, K
- E) No change

Ans: E

[15.22]. The data below refer to the following reaction:

$$2NO(g) + Br_2(g) \implies 2NOBr(g)$$

Concentration (M) [NO] [Br₂] [NOBr] 2.5 Initial 5.0 Equilibrium 2.0

Find the concentration of NOBr when the system reaches equilibrium.

Ans: 1.5 M

[15.23] If the reaction $2H_2S(q) \rightleftharpoons 2H_2(q) + S_2(q)$ is carried out at $1065^{\circ}C$, $K_p = 0.0120$. Starting from pure H2S introduced into an evacuated vessel at 1065°C, what will the total pressure in the vessel be at equilibrium if the equilibrated mixture contains 0.300 atm of $H_2(g)$? A) 1.06 atm B) 1.36 atm C) 2.39 atm D) 4.20 atm E) 1.51 atm Ans: E [15.24]. In the reaction: $2H_2O(1) \leftarrow H_3O^+(aq) + OH^-(aq)$ the conjugate acid-base pairs are A) *pair 1*: H₂O and H₂O; pair 2: H₃O⁺ and OH⁻ B) pair 1: H₃O⁺ and OH⁻; pair 2: H₃O⁺ and H₂O C) *pair 1*: H₃O⁺ and OH⁻; pair 2: OH and H₂O D) pair 1: H₂O and OH⁻; pair 2: H₂O and H₃O⁺ E) *pair 1*: H₃O⁺ and HO⁻: pair 2: OH and H₃O⁺ Ans: D [15.25]. What is the H⁺ ion concentration in a 4.8×10^{-2} M KOH solution? A) 4.8×10^{-2} M D) $4.8 \times 10^{-12} \text{ M}$ B) 1.0×10^{-7} M E) 2.1×10^{-13} M C) $4.8 \times 10^{-11} M$ Ans: E [15.26]. A 0.10 M HF solution is 8.4% ionized. Calculate the H⁺ ion concentration. D) 0.084 M E) 8.4×10^{-3} M C) 0.10 M A) 0.84 M B) 0.12 M

Ans: E

	A 5.5 L sample at is the pH of the		IO3 solution is mixed	with 1.2 L of a (0.34 M HCl solution.				
,	A) 0.23	B) 0.57	C) 1.07	D) 0.50	E) 0.84				
Ans: B									
[15.28].	Arrange the ac	ds H ₂ Se, H ₂ Te,	and H ₂ S in order of i	ncreasing acid	strength.				
	A) $H_2S < H_2Se < H_2Te$ B) $H_2S < H_2Se$ C) $H_2Te < H_2Se$ D) $H_2Se < H_2Se < H_2Te$ E) $H_2Se < H_2Te < H_2Se$								
Ans: A									
 [15.29]. When comparing acid strength of binary acids HX, as X varies within a particular group of the periodic table, which one of these factors dominates in affecting the acid strength? A) bond strength B) electron withdrawing effects C) percent ionic character of the H–X bond D) solubility E) Le Châtelier's principle Ans: A									
[15.30].	15.30]. Predict the direction in which the equilibrium will lie for the reaction H₂CO₃ + F⁻ ← HCO₃⁻ + HF.								
	$K_{a1}(H_2CO_3) = 4.2 \times 10^{-7}$; $K_a(HF) = 7.1 \times 10^{-4}$								
	A) to the right	B)	to the left	C) in the mide	lle				

Ans: B

[15.31]. Calculate the pH of a 0.055 M solution of CH₃COONa ($K_a(CH_3COOH) = 1.8 \times 10^{-5}$)

- A) 12.74
- B) 4.74
- C) 8.74
- D) 9.26
- E) 5.26

Ans: C

[15.32] The equilibrium constant for the reaction

$$C_7H_{15}COOH(aq) + HCOO^-(aq)$$
 \longrightarrow $C_7H_{15}COO^-(aq) + HCOOH(aq)$

is 7.23×10^{-2} at 25°C. If K_a for formic acid (HCOOH) is 1.77×10^{-4} , what is the acid dissociation constant for C₇H₁₅COOH?

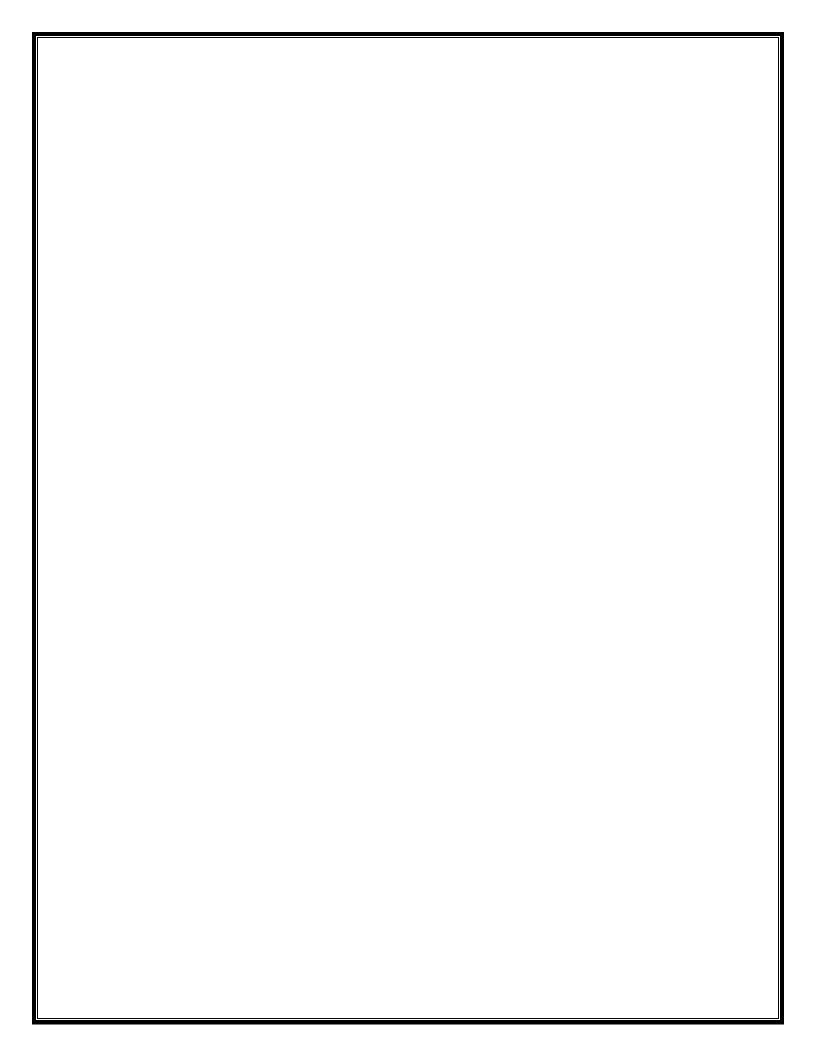
- A) 2.45×10^{-3}
- D) 1.00×10^{-4}
- B) 4.08×10^{-2}
- E) 1.28×10^{-5}
- C) 7.81×10^{-4}

Ans: E

[15.33] Which of these species will act as a Lewis acid?

A) NH₃ B) NH₄⁺ C) H₂O D) BF₃ E) F⁻

Ans: D



Chapter 16: Acid-Base Equilibria and Solubility Equilibria

A table of ionization constants and Ka's is required to work some of the problems in this chapter

- [16.1]. Which of the following yields a buffer solution when equal volumes of the two solutions are mixed?
 - A) 0.050 M H₃PO₄ and 0.050M HCl
 - B) 0.050M H₃PO₄ and 0.025 M HCl
 - C) 0.050M NaH₂PO₄ and 0.025M NaOH
 - D) 0.050M Na₃PO₄ and 0.050M M NaOH
 - E) 0.050M Na₃PO₄ and 0.025M NaOH

Ans: C

- [16.2] Which of the following is the most acidic solution?
 - A) 0.10 M CH₃COOH and 0.10 M CH₃COONa
 - B) 0.10 M CH₃COOH
 - C) 0.10 M HNO₂
 - D) 0.10 M HNO₂ and 0.10 M NaNO₂
 - E) 0.10 M CH₃COONa

Ans: C

- [16.3] You have 500.0 mL of a buffer solution containing 0.30 M acetic acid (CH₃COOH) and 0.20 M sodium acetate (CH₃COONa). What will the pH of this solution be after the addition of 20.0 mL of 1.00 M NaOH solution? [$K_a = 1.8 \times 10^{-5}$]
 - A) 4.65
- B) 4.71
- C) 4.56
- D) 4.84
- E) 5.07

Ans: B

	olution. After 25.00	•		10 ⁻⁴) is titrated wit , the pH in the titra				
	A) 2.17	B) 3.35	C) 2.41	D) 1.48	E) 7.00			
Ans: B	I							
[16.5]	The molar solution compound?	ubility of mangan	ese(II) carbonate	is 4.2 × 10 ⁻⁶ M. W	hat is K _{sp} for this			
Ans: D	,	B) 8.4×10^{-6}	C) 3.0×10^{-16}	D) 1.8 × 10 ⁻¹¹	E) 2.0 ×10 ⁻³			
 [16.6]	A saturated sodium carbonate solution at 100°C contains 45.5 g of dissolved sodium carbonate per 100. mL of solution. The solubility product constant for sodium carbonate at this temperature is							
Ans: E		0.316 C)	0.0790 D) 36.8 E)	316			
 [16.7]	Solid sodium iodide is slowly added to a solution that is 0.0050 M Pb $^{2+}$ and 0.0050 M Ag $^{+}$. [K _{sp} (PbI ₂) = 1.4 × 10 $^{-8}$; K _{sp} (AgI) = 8.3 × 10 $^{-17}$] Calculate the Ag $^{+}$ concentration when PbI ₂ just begins to precipitate.							
<mark>Ans: 5</mark>	$.0 \times 10^{-14} M$							
 [16.8]	NaCl is added slowly to a solution that is 0.010 M each in Cu+, Ag+, and Au+. The K _{sp} 's for CuCl, AgCl, and AuCl are 1.9×10^{-7} , 1.8×10^{-10} , and 2.0×10^{-13} , respectively.							
Which	compound will pr	ecipitate first?						
Ans: A	uCl							

	•		of a weak acid on the weak acid of	_	r in a solution contai	ning the salt NaA			
Ans: Fals	se								
					350. mL of a 0.150 ld3) = 1.8×10^{-5})	M solution of			
A)	7.6 g	B) 2.4 g	C) 5.4 g	D) 11 g	E) 3.3 g				
Ans: A									
	In which of the contraction of t		lowing solution	s will acetic a	cid have the greates	t percent			
	A) 0.1 M CH ₃ COOH B) 0.1 M CH ₃ COOH dissolved in 1.0 M HCI C) 0.1 M CH ₃ COOH plus 0.1 M CH ₃ COONa D) 0.1 M CH ₃ COOH plus 0.2 M CH ₃ COONa								
Ans: A									
[16.12].	Which o		g yields a buffe	er solution wh	en equal volumes of	the two solutions			
	B) 0.		and 0.15 M Nai		0M HClO4 and 0.10 N 5M HBr and 0.15M N				
Ans: C	; 								

[16.13].	[16.13]. Calculate the pH of a buffer solution that contains 0.25 M benzoic acid (C ₆ H ₅ CO ₂ H) an 0.15M sodium benzoate (C ₆ H ₅ COONa). [$K_a = 6.5 \times 10^{-5}$ for benzoic acid]							
			`	, -		cid]		
Ans: A	A) 3.9	37 B) 4.8	3 C) 4.19	D) 3.40	E) 4.41			
SC		a pH of 4.00 ±	•	•	c acid - sodium ac Ⅎ₃COOH to CH₃CC			
	A) 0.	18 B) 0.8	34 C) 1.19	D) 5.50	E) 0.10			
Ans: D	 Calculate th	 ne nH of a sol	ution that is 0.4	 410 M in HOCI	 and 0.050 M in Na	 OCI		
[10.13].	Calculate III	·			and 0.000 W III Ne	100i.		
			[Ka(HOCI) = 3]	.2 × 10 °]				
A)	0.39	B) 3.94	C) 6.58	D) 7.49	E) 8.40			
Ans: C								
[16.16].	For which	type of titration	on will the pH t	oe basic at the	equivalence point?)		
٨٠٠	B) S C) V	strong acid vs strong acid vs Veak acid vs.		D) All of th E) None o	e above. f the above.			
Ans	<mark>. U</mark>							

[16.17]. 24.00 mL of a 0.25 M NaOH solution is titrated with 0.10M HCl. What is the pH of the solution after 24.00 mL of the HCl has been added?

- A) 13.40
- B) 13.17
- C) 11.56
- D) 12.88
- E) 7.00

Ans: D

The solubility product for calcium phosphate is $K_{sp} = 1.3 \times 10^{-26}$. What is the molar [16.18]. solubility of calcium phosphate?

$$Ca_3 (PO_4)_2 \implies 3Ca^{2+} (aq) + 2PO_4^{3-} (aq)$$

- A) $1.3 \times 10^{-26} \,\text{M}$ D) $4.6 \times 10^{-6} \,\text{M}$
- B) 1.5×10^{-7} M
- E) $6.6 \times 10^{-6} \,\mathrm{M}$
- C) 2.6×10^{-6} M

Ans: C

[16.19]. A saturated sodium carbonate (Na₂CO₃) solution at 0°C contains 7.1 g of dissolved sodium carbonate per 100. mL of solution. The solubility product constant for sodium carbonate at this temperature is

- A) 1.2
- B) 0.30 C) 3.0×10^{-4}
 - D) 0.90
- E) 1.2×10^{-3}

Ans: A

[17.1]	Which of th	ese species wo	ould you expect	to have the low	est standard entropy (S°)
	A) Br ₂ (I)	B) Cl ₂ (g)	C) F ₂ (g)	D) H ₂ (g)	E) I ₂ (s)
Ans: E					
[17.2]	_	e following sub s), SiF4(g), SiH		order of increasi	ng entropy at 25°C.
			$lowest \rightarrow high$	hest	
Ans: C	,	B) HF(g) < A C) Al(s) < N D) Al(s) < F	Al(s) < NaF(s) < aF(s) < HF(g) < HF(g) < NaF(s)	(s) < HF(g) < Al(: SiF4(g) < SiH4(g) : SiH4(g) < SiF4(< SiF4(g) < SiH4 < SiF4(g) < SiH4	g) g) (g)
[17.3]	Which one	of the following	g reactions wou	ıld you expect to	have highest ∆S°?
Ans: D	I	B) C ₂ H ₂ C) C ₂ H ₂	$_{2}(g) + _{5/2}O_{2}(g){4}(g) + _{3}O_{2}(g) \rightarrow$	$CO_2(g) + 2H_2O(g)$ $\rightarrow 2CO_2(g) + H_2O(g)$ $\rightarrow 2CO_2(g) + 2H_2O(g)$ $\rightarrow 2CO_2(g) + 3H_2O(g)$) O(g) O(g)

[17.4] Sulfur can be separated from lead in the mineral galena, PbS(s), by "roasting" the ore in the presence of oxygen as shown in the following reaction:

$$2PbS(s) + 3O_2(g) \rightarrow 2PbO(s) + 2SO_2(g)$$

Calculate ΔS° for this reaction using the thermodynamic data provided below.

Compound	S°(J/K-mol)
PbS(s)	91.2
O ₂ (g)	205.0
PbO(s)	69.45
SO ₂ (g)	248.5

- A) –410 J/K·mol D) 21.8 J/K·mol B) –161.5 J/K·mol E) 43.5 J/K·mol
- C) -47.7 J/K·mol

Ans: B

- With respect to the system only, a reaction with $\Delta H > 0$ and $\Delta S < 0$ is predicted to be: [17.5]
 - A) Spontaneous at all temperatures
 - B) Spontaneous at high temperatures only
 - C) Spontaneous at low temperatures only
 - D) Nonspontaneous at all temperatures

Ans: D

- [17.6] At 1500°C the equilibrium constant for the reaction CO(g) + 2H₂(g) CH₃OH(g) has the value $K_p = 1.4 \times 10^{-7}$. Calculate ΔG° for this reaction at 1500°C.
 - A) 105 kJ/mol
- D) -105 kJ/mol
- B) 1.07 kJ/mol
- E) 233 kJ/mol
- C) -233 kJ/mol

Ans: E

[17.7] Calculate K_p at 298 K for the reaction $SO_2(g) + NO_2(g) \implies SO_3(g) + NO(g)$.

	ΔG^o f	
SO ₂ (g) SO ₃ (g) NO(g) NO ₂ (g)	–300.4 kJ/mol –370.4 kJ/mol 86.7 kJ/mol 51.8 kJ/mol	

A) 6.99×10^{-7} B) 5.71×10^{-8}

C) 14.2 D) 475

E) 1.42×10^6

Ans: E

[17.8] For the reaction SbCl₅(g) \Longrightarrow SbCl₃(g) + Cl₂(g),

 $\Delta G^{\circ}f$ (SbCl₅) = -334.34 kJ/mol

 $\Delta G^{\circ}f$ (SbCl₃) = -301.25 kJ/mol

 $\Delta H^{\circ}f$ (SbCl₅) = -394.34 kJ/mol

 $\Delta H^{\circ}f$ (SbCl₃) = -313.80 kJ/mol

Will this reaction proceed spontaneously at 298 K and 1 atm pressure?

Ans: No

[17.9] Consider the reaction $CO(g) + 2H_2(g) \leftarrow CH_3OH(I)$ at 25°C.

 $\Delta G^{\circ}f(CO) = -137.3 \text{ kJ/mol}$

 $\Delta G^{\circ}f$ (CH₃OH) = -166.3 kJ/mol

 $\Delta H^{\circ}_{f}(CO) = -110.5 \text{ kJ/mol}$

 $\Delta H^{\circ}f$ (CH₃OH) = -238.7 kJ/mol

 $S^{\circ}(CO) = 197.9 \text{ J/K-mol}$

 $S^{\circ}(CH_3OH) = 126.8 \text{ J/K-mol}$

Calculate value of the equilibrium constant (K_P) for this reaction at 25°C.

Ans: 1.21 × 10⁵

[17.10]	Given the following da	ta, calculate the	e boiling point of HC 	OOH (formic acid).	
		ΔH°_{f} (kJ/mol)	S° (J/K·mol)		
	HCOOH(I) –410 g) –363	251		
Ans: 115	S°C				
 [17.11]	Using the thermodynar	nic data provid		K _a for HCN(<i>aq</i>) at 2	5°.
		S° (
H ⁺ (<i>aq</i>)	0		0		
CN ⁻ (aq)			17.99 128.9 		
Ans: 2.8	× 10 ⁻⁹ (compared to the	e experimental	value of 4.9 × 10 ⁻¹⁰)	
 [17.12].	Which of these specie	s has the highe	est entropy (S°) at 25	5°C?	
	A) CO(g)	3) CH ₄ (g)	C) NaCl(s)	D) H ₂ O(I)	E) Fe(s)
Ans: B					

[17.13]. Aluminum forms a layer of aluminum oxide when exposed to air which protects the bulk metal from further corrosion.

$$4AI(s) + 3O_2(g) \rightarrow 2AI_2O_3(s)$$

Using the thermodynamic data provided below, calculate ΔS° for this reaction.

	S°(J/K⋅mol)	
Al(s) O ₂ (g) Al ₂ O ₃ (s)	28.3 205.0 50.99	

- A) 182.3 J/K⋅mol
- D) -626.2 J/K·mol
- B) 131.5 J/K·mol
- E) -802.9 J/K·mol
- C) -182.3 J/K·mol

Ans: D

[17.14]. Nitrosyl chloride (NOCI) decomposes at elevated temperatures according to the equation

 $2NOCl(g) \rightleftharpoons 2NO(g) + Cl_2(g)$. Calculate K_P for this reaction at 227°C.

 $(\Delta H^{\circ} = 81.2 \text{kJ/mol}, \Delta S^{\circ} = 128 \text{ J/K·mol})$

- A) 1.59×10^{-2} B) 2.10×10^{-7} C) 62.8 D) 4.90×10^{6} E) 3.20×10^{9}

Ans: A

[17.15] In the gas phase, methyl isocyanate (CH₃NC) isomerizes to acetonitrile (CH₃CN), $H_3C-N=C(g)$ \longleftrightarrow $H_3C-C=N(g)$

with $\Delta H^{\circ} = -89.5$ kJ/mol and $\Delta G^{\circ} = -73.8$ kJ/mol at 25°C. Find the equilibrium constant for this reaction at 100°C.

A) 1.68×10^{-10}

D) 4.63×10^{-11}

B) 5.96×10^9

E) 8.64×10^{12}

C) 2.16×10^{10}

[17.16]	Assuming ΔS° and	∆H° do not var	with temperature,	at what tempera	ture will the
rea	ction shown below	become spontar	neous?		

$$C(s) + H_2O(g) \rightarrow H_2(g) + CO(s)$$
 ($\Delta S^{\circ} = 133.6 \text{ J/K} \cdot \text{mol}$; $\Delta H^{\circ} = 131.3 \text{ kJ/mol}$)

Ans: 710°C

[17.17].

At 700 K, the equilibrium constant for the reaction $CO(g) + H_2O(g)$ $CO_2(g) + H_2(g)$ is 5.10. What is ΔG° for this reaction at this temperature?

Ans: -9.48 kJ/mol

Chapter 18: Electrochemistry NOTE: A table of standard reduction potentials is required to work many of these problems. [18.1] What is the total number of moles (n) of electrons exchanged between the oxidizing agent and the reducing agent in the overall redox equation: $5 \text{ Ag}^+ (aq) + \text{Mn}^{2+} (aq) + 4\text{H}_2\text{O}(l) \rightarrow 5\text{Ag}(s) + \text{MnO}_4^- (aq) + 8\text{H}^+ (aq)$ B) 2 A) 1 C) 3 D) 5 E) 7 Ans: D [18.2] Complete and balance the following redox equation. What is the coefficient of H2O when the equation is balanced with the set of smallest whole-number coefficients? $H_2O + MnO_4^- + I^- \rightarrow MnO_2 + IO_3^-$ (basic solution) A) 1 B) 2 C) 4 D) 10 E) None of these. Ans: A [18.3] Given the following cell diagram, $Al(s) | Al^{3+}(aq) | | Cd^{2+}(aq) | Cd (s)$ what is the balanced overall (net) cell reaction? A) Al(s) + $Cd^{2+}(aq) \rightarrow Al^{3+}(aq) + Cd$ (s) B) $Al^{3+}(aq) + Cd(s) \rightarrow Al(s) + Cd^{2+}(aq)$ C) $2AI(s) + 3Cd^{2+}(aq) \rightarrow 2AI^{3+}(aq) + 3Cd(s)$ D) $2Al^{3+}(aq) + 3Cd(s) \rightarrow 2Al(s) + 3Cd^{2+}(aq)$ E) $3AI^{3+}(aq) + 2Cd(s) \rightarrow 3AI(s) + 2Cd^{2+}(aq)$ Ans: C

[18.4] A galvanic cell has the overall reaction:

$$2Fe(NO_3)_2(aq) + Pb(NO_3)_2(aq) \rightarrow 2Fe(NO_3)_3(aq) + Pb(s)$$

Which is the half reaction occurring at the cathode?

A)
$$NO^{3-}$$
 (aq) + $4H^{+}$ (aq) + $3e_{-} \rightarrow NO(g) + 2H_{2}O(I)$

B)
$$NO^{3-}(aq) + 2H^{+}(aq) + e_{-} \rightarrow NO_{2}(g) + H_{2}O(I)$$

C)
$$Fe^{2+}$$
 (aq) $\to Fe^{3+}$ (aq) + e-

D)
$$Pb^{2+}(aq) + 2e \rightarrow Pb(s)$$

D)
$$Pb^{2+}$$
 (aq) + 2e \rightarrow Pb(s)
E) Fe^{2+} (aq) + e \rightarrow Fe³⁺ (aq)

Ans: D

[18.5] Calculate E°cell for a silver-aluminum cell in which the cell reaction is

$$Al(s) + 3Ag^{+}(aq) \rightarrow Al^{3+}(aq) + 3Ag(s)$$

- A) -2.46 V
- B) 0.86 V
- C) -0.86 V
- D) 2.46 V
- E) none of these

Ans: D

Consider the following electrochemical cell: [18.6]

If the standard cell emf is 3.16 V, what is the standard reduction potential for uranium?

- A) -3.16 V
- B) +3.16 V
- C) -1.80 V
- D) +1.80 V
- E) +1.36 V

Ans: C

[18.7] Consider the following standard reduction potentials in acid solution:

	E°(V)	
$Al^{3+} + 3e_{-} \rightarrow Al(s)$ $AgBr(s) + e_{-} \rightarrow Ag(s) + Br^{-}$ $Sn^{4+} + 2e_{-} \rightarrow Sn^{2+}$ $Fe^{3+} + e_{-} \rightarrow Fe^{2+}$	-1.66 +0.07 +0.14 +0.77	

The strongest oxidizing agent among those shown above is

- A) Fe^{3+} . B) Fe^{2+} . C) Br^{-} .
- D) Al³⁺.
- E) Al.

Ans: A

For the electrochemical cell Pt(s) | H₂(1 atm) | H⁺ (1 M) || Cu²⁺ (1 M) | Cu(s), which one of [18.8] the following changes will cause an increase in the cell voltage?

- A) Lower the H₂(g) pressure.
- B) Increase the size/mass of the copper electrode.
- C) Lower the H⁺ (aq) concentration.
- D) Decrease the concentration of Cu²⁺ ion.
- E) None of the above.

Ans: C

[18.9] Consider an electrochemical cell involving the overall reaction

$$2AgBr(s) + Pb(s) \rightarrow Pb^{2+} + 2Ag(s) + 2Br^{-}$$

Each half-reaction is carried out in a separate compartment. The anion included in the lead half-cell is NO₃⁻. The cation in the silver half-cell is K⁺. The two half-cells are connected by a KNO₃ salt bridge. If [Pb²⁺] = 1.0 M, what concentration of Br⁻ ion will produce a cell emf of 0.25 V at 298 K?

Given: AgBr(s) +
$$e_- \rightarrow Ag + Br^-$$
, $E^{\circ} = +0.07 \text{ V}$.

A) 0.02 M

B) 0.14 M C) 0.38 M D) 1.0 M E) 7.0 M

_	-	oulombs (C) of ele Ca ²⁺ ion to calciu	ectrical charge mu um metal?	ust pass throu	gh an electro	olytic cell to
Ans: B	A) 190,000 C	B) 85,000 C	C) 21,000) C D) 4	2,500 C	E) 0.88 C
		it take to produce ith a current of 2	e 78 g of Al metal .0 A?	by the reduct	tion of Al ³⁺ in	an
	A) 0.01 s	B) 420 s	C) 13 h	D) 116 h	E) 1.0 × 1	0 ¹² s
Ans: D						
[18.12]	Which eleme	nt is associated v	with the term "gal	vanized"?		
Ans: B	A) Ga	B) Zn C) (Cd D) Hg	E) Pb		
[18.13]			oles (<i>n</i>) of electron overall redox equ	_	between the	e oxidizing agent
	3C ₂ O ₄ ²⁻ (ad	q) + Cr ₂ O ₇ ²⁻ (aq)	$+ 14H^+(aq) \rightarrow 6$	CO ₂ (g) + 2Cr ³	ⁱ⁺ (aq) + 7H ₂ C	D(I)
Ans: 6						
	ectric current of	• •	e deposited on the state of the		•	
Ans: 0.7	<mark>51 g</mark>					

[18.15] Many different ways have been proposed to make batteries. One cell is set up with copper and lead electrodes in contact with CuSO₄(aq) and Pb(NO₃)₂ (aq), respectively. If the Pb²⁺ and Cu²⁺ concentrations are each 1.0 M, what is the overall cell potential?

$$Pb^{2+} + 2e^{-} \rightarrow Pb$$
 $E^{\circ} = -0.22 \text{ V}$ $Cu^{2+} + 2e^{-} \rightarrow Cu$ $E^{\circ} = +0.34 \text{ V}$

Ans: 0.56 V

[18.16] The electrochemical cell that utilizes the reaction $Zn + Cu^{2+}$ (1 M) $\rightarrow Zn^{2+}$ (1 M) + Cu will have a lower cell emf when the concentrations of Zn^{2+} and Cu^{2+} ions are decreased to 0.1 M.

Ans: False

[18.17] A metal object is to be gold-plated by an electrolytic procedure using aqueous AuCl₃ electrolyte. Calculate the number of moles of gold deposited in 3.0 min by a constant current of 10. A.

A)
$$6.2 \times 10^{-3}$$
 mol

D)
$$3.5 \times 10^{-5}$$
 mol

B)
$$9.3 \times 10^{-3}$$
 mol

C)
$$1.8 \times 10^{-2} \text{ mol}$$

Ans: A

[18.18]. What is the total number of moles (*n*) of electrons exchanged between the oxidizing agent and the reducing agent in the overall redox equation:

$$Cr_2O_7^{2-}(aq) + 6Fe^{2+}(aq) + 14H^+(aq) \rightarrow 2Cr^{3+}(aq) + 6Fe^{3+}(aq) + 7H_2O(l)$$

- A) 1
- B) 2
- C) 3
- D) 6
- E) 12

Ans: D

[18.19]. Consider the following standard reduction potentials in acid solution:

	E°(V)
$Al^{3+} + 3e^- \rightarrow Al(s)$	– 1.66
$AgBr(s) + e^{-} \rightarrow Ag(s) + Br^{-}$	+0.07
$\operatorname{Sn}^{4+} + 2e^{-} \rightarrow \operatorname{Sn}^{2+}$	+0.14
$Fe^{3+} + e^{-} \rightarrow Fe^{2+}$	+0.77

The strongest reducing agent among those shown above is

A) Fe^{3+} . B) Fe^{2+} . C) Br^{-} . D) Al^{3+} .

E) Al.

Ans: E

[18.20]. Consider a Galvanic cell constructed from the following half cells, linked by an external circuit and by a KCl salt bridge.

- an Al(s) electrode in 1.0 M Al(NO₃)₃ solution
- a Pb(s) electrode in 1.0 M Pb(NO₃)₂ solution

The balanced overall (net) cell reaction is

- A) $Pb(s) + Al^{3+}(aq) \rightarrow Pb^{2+}(aq) + Al(s.)$
- B) $3Pb(s) + 2Al^{3+}(aq) \rightarrow 3Pb^{2+}(aq) + 2Al(s)$.
- C) $3Pb^{2+}(aq) + 2AI(s) \rightarrow 3Pb(s) + 2AI^{3+}(aq)$.
- D) $Pb^{2+}(aq) + Al(s) \rightarrow Pb(s) + Al^{3+}(aq)$.

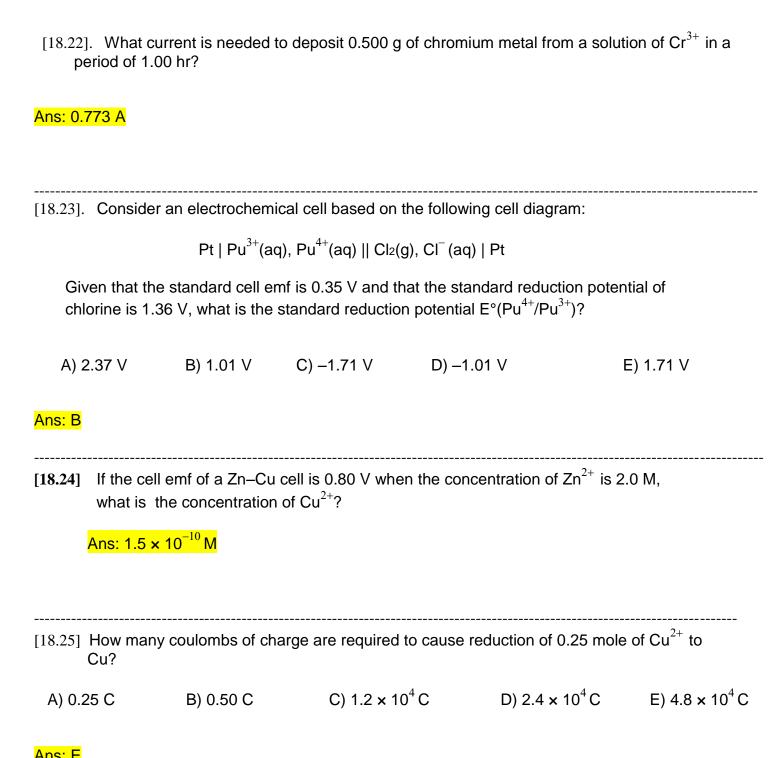
Ans: C

[18.21]

For the reaction $Ni^{2+}(aq) + 2Fe^{2+}(aq) \rightarrow Ni(s) + 2Fe^{3+}(aq)$, the standard cell potential E°cell is

A) +2.81 V. B) +1.02 V. C) +0.52 V. D) -1.02 V. E) -2.81 V.

Ans: D



[18.26] Complete and balance the following redox equation. What is the coefficient of H₂S when the equation is balanced using the set of smallest whole-number coefficients?

$$H_2S + MnO_4^- \rightarrow Mn^{2+} + SO_4^{2-}$$
 (acidic solution)

- A) 1
- B) 2 C) 4
- D) 5 E) None of these.

Ans: D

[18.27] Complete and balance the following redox equation. What is the coefficient of OH⁻ when the equation is balanced using the set of smallest whole-number coefficients?

 $MnO_4^- + \Gamma \rightarrow MnO_2 + IO_3^-$ (basic solution)

- A) 1 B) 2 C) 4
- D) 10
- E) None of these.

Chapter 19: Nuclear Chemistry

[19.1] 1. Alpha particles are identical to

- A) protons.
- D) helium nuclei.
- B) helium atoms.
- E) electrons.
- C) hydrogen atoms.

Ans: D

[19.2] When atoms of beryllium-9 are bombarded with alpha particles, neutrons are produced. What new isotope is also formed?

$$\frac{4}{2}$$
He + $\frac{9}{4}$ Be $\to \frac{1}{0}$ n + _____

- A) ${}_{6}^{12}$ C B) ${}_{3}^{5}$ Li C) ${}_{3}^{8}$ Li D) ${}_{5}^{10}$ B E) ${}_{5}^{12}$ B

Ans: A

[19.3] Consider the following decay series:

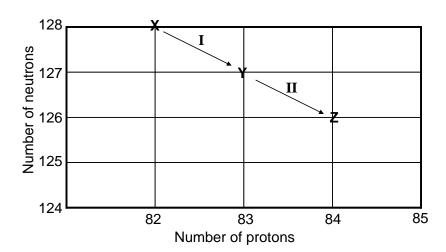
$$\overset{\scriptscriptstyle{218}}{\underset{\scriptscriptstyle{84}}{\text{Po}}} \text{Po} \overset{a}{\longrightarrow} \overset{\scriptscriptstyle{214}}{\underset{\scriptscriptstyle{82}}{\text{Pb}}} \text{Pb} \overset{b}{\longrightarrow} \overset{\scriptscriptstyle{214}}{\underset{\scriptscriptstyle{83}}{\text{Bi}}} \overset{\text{C}}{\longrightarrow} \overset{\scriptscriptstyle{214}}{\underset{\scriptscriptstyle{84}}{\text{Po}}} \text{Po} \overset{d}{\longrightarrow} \overset{\scriptscriptstyle{210}}{\underset{\scriptscriptstyle{82}}{\text{Pb}}} \text{Pb}$$

Which type of nuclear process occurs in steps c and d, respectively?

- A) α emission, α emission
- D) β emission, α emission
- α emission, β emission B)
- E) positron emission, β emission
- C) β emission, positron emission

Ans: D

[19.4] Consider the following decay series:

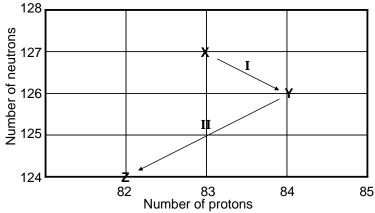


What is the symbol for the product labeled as Z?

- A) 127 Bi
- B) 128 Pb
- C) 210 Bi
- D) 210 Pb
- E) ²¹⁰₈₄Po

Ans: E

[19.5] Consider the following decay series:



What type of nuclear process occurs at the transformation labeled I?

A) α emission

D) electron capture

B) β emission

E) gamma radiation

C) positron emission

A balanced nuclear equation representing the beta emission of iodine-131 is which of the following? [19.6]

- 53 l A) 131
- B)
- 131 C) 53
- 131 53 131 D)
- E)

Ans: A

[19.7] Which of the following nuclear processes results in an increase of atomic number in the product element formed?

- Alpha emission I.
- Beta emission II.
- III. Positron emission
- IV. Electron capture
- A) I and II B) III and IV C) I, II, and III D) II only E) IV only

Ans: D

The energy equivalent of a mass defect of -0.1710 amu is which of the following? [19.8]

 $-5.130 \times 10^7 \text{ J}$ A)

D)

 $-1.539 \times 10^{16} \mathrm{J}$ B)

 $-2.556 \times 10^{-11} \text{ J}$ $-8.519 \times 10^{-20} \text{ J}$ E)

 $-2.556 \times 10^{-8} \text{ J}$

Ans: D

[19.9] Find the nuclear binding energy of potassium-40 (atomic mass = 39.9632591 amu) in units of joules per nucleon. [Data: neutron mass = 1.674928×10^{-24} g; proton mass = 1.672623×10^{-24} g; electron mass = 9.109387×10^{-28} g; $N_A = 6.0221367 \times 10^{23}$ /mol; $c = 2.99792458 \times 10^{8}$ m/s]

- A)
- D)
- $\begin{array}{l} 1.37\times 10^{-12}~\text{J/nucleon} \\ 5.48\times 10^{-11}~\text{J/nucleon} \end{array}$ B)
- 1.41×10^{-12} J/nucleon 2.97×10^{-12} J/nucleon E)
- 5.64×10^{-11} J/nucleon C)

Ans: A

Calculate the energy released in joules when one mole of polonium-214 decays according [19.10] to the equation

$$\frac{214}{84}$$
 Po $\to \frac{210}{82}$ Pb + $\frac{4}{2}$ He.

[Atomic masses: Pb-210 = 209.98284 amu, Po-214 = 213.99519 amu, He-4 = 4.00260 amu.]

 $8.78\times10^{14}\ J/mol$ A)

D) $-9.75 \times 10^{-3} \text{ J/mol}$ E) $1.46 \times 10^{-9} \text{ J/mol}$

 $7.2 \times 10^{14} \text{ J/mol}$ $8.78 \times 10^{11} \text{ J/mol}$ B)

Ans: C

- [19.11] Polonium-208 is an alpha emitter with a half-life of 2.90 years. How many milligrams of polonium from an original sample of 2.00 mg will remain after 8.00 years?
 - A) 0.147 mg B) 0.296 mg C) 0.725 mg D) 6.77 mg E) 1.90 mg

Ans: B

- [19.12] A rock contains 0.37 mg of Pb-206 and 0.95 mg of U-238. The half-life of the decay series U-238 \rightarrow Pb-206 is 4.5×10^9 yr. Assuming no Pb-206 was present in the rock initially, how old is the rock?
 - $1.7 \times 10^9 \text{ yr}$ A)

D) $4.5 \times 10^9 \text{ yr}$ E) $2.4 \times 10^9 \text{ yr}$

 $5.2 \times 10^9 \text{ yr}$ $2.7 \times 10^6 \text{ yr}$ B)

- - Ans: E
- [19.13] The Rb-87/Sr-87 method of dating rocks is often used by geologists:

$$^{87}_{37}\text{Rb} \rightarrow ^{87}_{38}\text{Sr} + ^{0}_{-1}\beta \qquad t_{1/2} = 6.0 \times 10^{10}\,\text{yr}$$

Estimate the age of a rock sample in which the present-day mole ratio of Rb-87 to Sr-87 is 36:1.

 $2.4 \times 10^{9} \text{ yr}$ A)

D) $4.1 \times 10^{-11} \text{ yr}$ E) $3.6 \times 10^{11} \text{ yr}$

 $1.7 \times 10^9 \text{ yr}$ $3.1 \times 10^{11} \text{ yr}$ B)

Ans: A

[19.14] Decay of ruthenium-93 by positron emission yields

A) technetium-93.

D) rhenium-93.

B) molybdenum-89.

E) ruthenium-92.

C) palladium-97.

Ans: A

[19.15] Strontium-90 has a half-life of 28.8 years. How much strontium-90 was present initially, if after 144 years 10.0 g remain?

Ans: 320 g

[19.16] Balance the equation ${}^{235}_{92}U + {}^{1}_{0}n \rightarrow ? + {}^{97}_{40}Zr + 2{}^{1}_{0}n$.

Ans: ${}^{235}_{92}U + {}^{1}_{0}n \rightarrow {}^{137}_{52}Te + {}^{97}_{40}Zr + 2{}^{1}_{0}n$

[19.17] The isotope of iodine with mass number 128 undergoes beta decay with a half-life of 25.00 minutes. How long will it take for 93.75% of a sample of pure ¹²⁸ I to decay?

Ans: 100. min

Chapter 24. Organic Chemistry

[24.1] Alkanes have the general formula

- A) C_nH_{2n-4}
- B) C_nH_{2n-2}
 - C) C_nH_{2n}
- D) C_nH_{2n+2}
- E) C_nH_{2n+4}

Ans: D

[24.2] Cycloalkanes have the general formula

- A) C_nH_{2n-4}
- B) C_nH_{2n-2}
 - C) C_nH_{2n}
- D) C_nH_{2n+2}
- E) C_nH_{2n+4}

Ans: C

[24.3] Select the one molecule that is a *saturated hydrocarbon*.

- A) C₂H₄
- B) C₄H₈
- C) C₄H₆
- D) C₄H₁₀
- E) C_4H_2

Ans: D

[24.4] Which one of these hydrocarbons does *not* have isomers?

- A) C₇H₁₆
- B) C₆H₁₄
- C) C₅H₁₀
- D) C₄H₈
- E) C₃H₈

Ans: E

Which of these pairs are geometric isomers? [24.5]

> A) CH₃CH₂-O-CH₂CH₃ and CH₃CH₂CH₂CH₂OH

> B) C1 CH3-CH-CH3 and ClCH2CH2CH2CH2Cl

- C) C1 $CH_3CH_2C=CH$ and $CH_3-C=\overset{\downarrow}{C}-CH_2C1$ $\overset{\downarrow}{C1}$ $\overset{\downarrow}{C1}$ $\overset{\downarrow}{H}$
- D) Η ClCH2-C=C-H and ClCH2-C=C-H H CH2C1

Ans: D

[24.6] The molecule CHBr₂CHCl₂ has how many chiral centers?

- A) 0
- B) 1
- C) 2
- D) 3
- E) 4

Ans: A

[24.7] Which option represents dimethyl ether?

> A) CH₃CH₂OH

CH₃OCH₃ D)

B) CH₃OH

CH₃CH₂OCH₃ E)

C) CH₃COCH₃

Ans: D

Which one of the following functional groups is found in *alcohols*. [24.8]

- A. $R-\overset{\bullet}{\bigcirc}-H$ B. $R-\overset{\bullet}{\bigcirc}-R'$ C. $R-\overset{\bullet}{\bigcirc}-R'$ D. $R-\overset{\bullet}{\bigcirc}-H$ E. $R-\overset{\bullet}{\bigcirc}-R'$

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

Ans: A

- [24.9] The name for the compound with the formula CH₃CH₂CH₂CH₂OH is
 - A) propanol. B) propane. C) butanol. D) pentane. E) pentanol.

Ans: C

- [24.10] The expected product from the addition of HCl to CH₃-CH₂-CH=CH₂ is
 - A) CH₃-CH₂-CH=CHCl
- D) CH₃-CH₂-CH₂-CH₂Cl
- B) $CH_3-CH_2-CCl=CH_2$

- E) CH₃-CH₂-CHCl-CH₃
- C) CH₃-CHCl-CH=CH₂

Ans: E

- [24.11] Bromination of benzene (C₆H₆), an aromatic compound,
 - A) occurs by substitution rather than addition.
 - B) occurs by addition rather than substitution.
 - C) occurs more rapidly than bromination of a nonaromatic compound.
 - D) results in formation of 1,2,3,4,5,6-hexabromocyclohexane.
 - E) occurs in the absence of a catalyst.

Ans: A

[24.12] Name the following compound:

$$\begin{array}{c} CH_3 \\ | \\ CH_3-CH-CH-CH_3 \\ | \\ CH_3 \end{array}$$

Ans: 2,3-dimethylbutane

[24.13] Write the formula for the alcohol and the carboxylic acid from which the following ester may be synthesized.

Ans:

$$\rm CH_3-CH_2-COOH$$
 and $\rm CH_3-CH-CH_2-OH$ $|$ $\rm CH_3$

- [24.14] Which type of organic compound does *not* contain a carbonyl group?
 - A) ethers
- B) carboxylic acids
- C) ketones
- D) aldehydes
- E) esters

Ans: A

[24.15] Water and ethylene are combined at high temperature and pressures in the presence of sulfuric acid to produce ethanol according to the following reaction:

$$CH_2=CH_2(g) + H_2O(g) \rightarrow CH_3CH_2OH(g)$$

What is the hybridization of the bolded carbon atoms in the reactants and products, respectively? A) $sp^3 sp^2$ B) $sp^2 sp^3$ C) $sp sp^2$ D) $sp^3 sp$ E) $sp^2 sp^2$

CHEM 1412 Formulas and Constants

 $mass\ of\ proton\ =\ 1.00728\ amu$

 $c = 3.00 \times 10^8 \text{ m/s}$

 $K = {}^{0}C + 273.15$ $1 g = 6.022 \times 10^{23} \text{ amu}$ $mass\ of\ neutron\ =\ 1.00866\ amu$

 $F = 96500 \text{ C/(mol of e^{-})} = 96500 \text{ J/(V mol of e^{-})}$

R = 0.08206 (L atm)/(mol K) = 8.314 J/(mol K)

1 atm = 760 mm Hg

$$S_g = kP_g$$

$$\mathbf{P_A} = \mathbf{X_A} \mathbf{P_A}^{\mathbf{o}}$$

$$\Delta T_f = K_f m$$

$$\Delta T_b = K_b m$$

$$\pi = MRT$$

$$PV = nRT$$

$$[A]_t = -kt + [A]_0$$
 $t_{1/2} = [A]_0 / 2k$

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$$\ln\left([\mathbf{A}]_{t}/[\mathbf{A}]_{0}\right) = -\mathbf{k}t$$

$$ln[A]_t = -kt + ln[A]_0$$
 $t_{1/2} = 0.693 / k$

$$t_{1/2} = 0.693 / k$$

$$1/[A]_t = kt + 1/[A]_0$$
 $t_{1/2} = 1/(k[A]_0)$

$$t_{1/2} = 1/(k[A]_0)$$

$$k = Ae^{-Ea/RT}$$

$$ln(k_1/k_2) = E_a/R (1/T_2 - 1/T_1)$$

For the chemical reaction (at Equilibrium):

$$aA + bB \rightleftharpoons cC + dD$$

$$Kc = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$
 (Law of mass action. General expression of equilibrium constant)

$$Kp=Kc~(0.0821*T)^{\Delta n}~$$
 , Relationship between Kp and Kc.
$$\Delta n=moles~of~gaseous~products-moles~of~gaseous~reactants.$$

$$Kc = Kc' \times Kc''$$
, The equilibrium constant for the overall reaction is given by the product of the equilibrium constants for the individual reactions.

Reaction Quotient (Qc) when instead of Equilibrium concentrations, the initial concentrations are used in the equilibrium constant expression. Qc determines the direction in which the net reaction will proceed:

(Qc < Kc: system proceeds from left to right, consuming reactants, forming products)

(Qc = Kc: system is at equilibrium)

(Qc > Kc: system proceeds from right to left, consuming products, forming reactants).

$$\begin{split} K_w &= [H^+] [OH^-] \\ pH &= -\log [H^+], \\ [H^+] &= 10^{-pH} \\ pOH &= -\log [OH^-], \\ [OH^-] &= 10^{-pOH} \\ pH &+ pOH &= 14 \end{split}$$

Ion-product constant of water.

Definition of pH of a solution.

Calculating H+ ion concentration from pH.

Definition of pOH of a solution.

Calculating OH ion concentration from pOH.

$$[H^{+}][OH^{-}] = 1.0 \times 10^{-14}$$

$$K_a K_b = K_w = 1.0 \times 10^{-14}$$

Relationship between the acid and base ionization constants of a conjugate acid-base pair.

Calculating pH of a solution containing a weak acid and a soluble salt of a weak acid (e.g. NaA):

$$HA(aq) + H2O(l) \rightarrow H3O^{+}(aq) + A^{-}(aq)$$

or

$$HA(aq) + H2O(l) \rightarrow H^{+}(aq) + A^{-}(aq)$$

The ionization constant Ka:

$$\mathbf{K_a} = \underline{[\mathbf{H}^+] [\mathbf{A}^-]}$$

$$pK_a = - \log K_a$$
,

Henderson-Hasselbach equation:

$$pH = pK_a + log \underline{[conjugate base]}$$
[conjugate acid]

Percent ionization = <u>ionized acid concentration at equilibrium</u> × 100% initial concentration of acid

Ka = acid ionization (dissociation) constant (measure of a strength of an acid)

$$Ka = \underline{[H3O^{+}][A^{-}]}$$

$$[HA]$$

Kb = base ionization (dissociation) constant (measure of a strength of a base).

$$\mathbf{Kb} = \underline{\mathbf{[BH}^{+}]} \underline{\mathbf{[OH}^{-}]}$$

$$\underline{\mathbf{[B]}}$$

The net direction of an acid-base reaction proceeds to the greater extent in the direction in which a stronger acid and stronger base form a weaker acid and weaker base.

Quadratic equation:
$$a x^2 + b x + c = 0$$

$$x = \frac{-b \pm SQRT(b^2 - 4 a c)}{2a}$$

$$pH = pK_a + log([base]/[acid])$$

$$\Delta G^o \,=\, \Delta H^o \,\,\textbf{-}\,\, T \, \Delta S^o \qquad \qquad \Delta G^o \,=\, \textbf{-RT ln } K_{eq}$$

$$\Delta G = \Delta G^{o} + RT \ln Q$$
 $\Delta G^{o} = -n F E^{o}$ $E = E^{o} - (0.0592/n) \log Q$

$$\Delta E = \Delta mc^2$$
 Coulombs = Amps X seconds

Soluble Compounds	Important Exceptions
Compounds containing NO ₃	none
Compounds containing C ₂ H ₃ O ₂ ⁻	none
Compounds containing Cl ⁻	Salts of Ag ⁺ , Hg ₂ ²⁺ , Pb ²⁺
Compounds containing Br	Salts of Ag ⁺ , Hg ₂ ²⁺ , Pb ²⁺
Compounds containing ☐	Salts of Ag ⁺ , Hg ₂ ²⁺ , Pb ²⁺
Compounds containing SO_4^{2-}	Salts of Ca ²⁺ , Sr ²⁺ , Ba ²⁺ , Hg ₂ ²⁺ , Pb ²⁺
Insoluble compounds	Important Exceptions
Compounds containing S ²⁻	Salts of ammonium, alkali metal cations and Ca ²⁺ , Sr ²⁺ , Ba ²⁺
Compounds containing CO ₃ ²⁻	Salts of ammonium, alkali metal cations
Compounds containing PO ₄ ³⁻	Salts of ammonium, alkali metal cations
Compounds containing OH	Salts of ammonium, alkali metal cations and Ca ²⁺ , Sr ²⁺ , Ba ²⁺

$$\begin{array}{ccc} \textbf{Percent by mass} & = & \underline{mass \ of \ solute} & \times \ 100\% \\ & & \underline{[mass \ of \ solute \ + \ mass \ of \ solvent]} \end{array}$$

Two-component solution:		
Mole fraction of solute A: X_A	=	moles of solute A
		moles of solute A + moles of solvent B
Molarity = number of moles of se	olute in 1 L o	f solution:
Molarity =		moles of solute
		Liters of solution
Molality = number of moles of so	olute dissolve	d in 1 kg (1000 g) of solvent:
Molality =		moles of solute
		Mass of solvent (1 kg)
Density = Mass / Volume.		