

Chemistry 1020, Module 15A

■ Review of Acids and Bases from Previous Modules (continued)

1b) For each of the following substances, write a balanced equation showing how the substance dissociates when placed in water. Then use the magnitude of the equilibrium expression to indicate the relative amounts of each of the species in the balanced equation.

S.	acetic acid	-Balanced equation for dissociation in water:	$\text{CH}_3\text{COOH}(\text{aq}) + \text{H}_2\text{O} \rightleftharpoons \text{CH}_3\text{COO}^-(\text{aq}) + \text{H}_3\text{O}^+(\text{aq})$
		-Relative amounts each species:	$K_a = 1.8 \cdot 10^{-5} \ll 1$ Therefore, much more CH_3COOH than CH_3COO^- or H_3O^+ in the solution at equilibrium.
A.	hydrofluoric acid	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	
B.	nitric acid	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	
C.	ammonia	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	
D.	hydrocyanic acid	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	
E.	CH_3NH_2	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	
F.	propionic acid	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	
G.	hydrochloric acid	-Balanced equation for dissociation in water:	
		-Relative amounts each species:	

✓✓✓■ **Calculation of the pH or pOH of a Solution of a Weak Acid or Base AGAIN**

2a) Work the indicated problem using an INITIAL/CHANGE/FINAL chart. (This is a review from Module 14B.)

S. What is the pH of a 0.52 M solution of acetic acid?

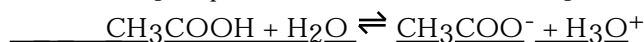
•Formula: CH₃COOH

•Type cmpd: acid

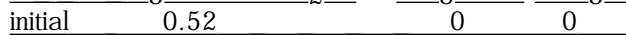
•Strength: weak

Prediction of pH: pH < 7

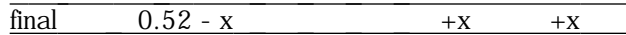
Reason for prediction: CH₃COOH is an acid.



$$x^2 = 0.52 \cdot (1.8 \cdot 10^{-5}) = 9.36 \cdot 10^{-6}$$



$$x = 3.06 \cdot 10^{-3}$$



K_a for acetic acid = 1.8 · 10⁻⁵

$$\text{Check: } \frac{x}{0.52} = \frac{3.06 \cdot 10^{-3}}{0.52} = 0.0059$$

$$K_a = \frac{[\text{CH}_3\text{COO}^-][\text{H}_3\text{O}^+]}{[\text{CH}_3\text{COOH}]} = 1.8 \cdot 10^{-5} = \frac{x \cdot x}{0.52 - x}$$

< 0.05 so ok

Assume 0.52 >> x so 0.52 - x = 0.52

Therefore, [H⁺] = 3.06 · 10⁻³ M

Therefore, equation reduces to $1.8 \cdot 10^{-5} = \frac{x \cdot x}{0.52}$

$$\text{pH} = -\log[\text{H}^+] = -\log(3.06 \cdot 10^{-3}) = 2.51$$

Agree with prediction: pH < 7 as predicted

A. What is the pOH of a 0.19 M solution of benzoic acid?

•Formula:

•Type cmpd:

•Strength:

Prediction of pOH:

Reason for prediction:

Agree with prediction:

B. What is the pH of a 0.26 M solution of hydrocyanic acid?

•Formula:

•Type cmpd:

•Strength:

Prediction of pH:

Reason for prediction:

Agree with prediction:

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✓✓✓■ *Calculation of the pH or pOH of a Solution of a Weak Acid or Base AGAIN (continued)*

2a) Work the indicated problem using an INITIAL/CHANGE/FINAL chart.

C. What is the pOH of a 0.53 M solution of formic acid?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pOH:

.....
Reason for prediction:

.....
Agree with prediction:

D. What is the pH of a 0.34 M solution of nitrous acid?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pH:

.....
Reason for prediction:

.....
Agree with prediction:

E. What is the pH of a 0.45 M solution of hydrofluoric acid?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pH:

.....
Reason for prediction:

.....
Agree with prediction:

✓✓✓■ Calculation of the pH or pOH of a Solution of a Weak Acid or Base AGAIN (cont'd)

2b) S. What is the pH of a 0.26 M solution of ammonia?

•Formula: NH₃

•Type cmpd: base

•Strength: weak

Prediction of pH: pH > 7

Reason for prediction: NH₃ is a base.

$$\begin{array}{cccc} & \text{NH}_3 + \text{H}_2\text{O} & \rightleftharpoons & \text{NH}_4^+ + \text{OH}^- \\ \text{initial} & 0.26 & & 0 \quad 0 \\ & -x & & +x \quad +x \\ \text{final} & 0.26 - x & & +x \quad +x \end{array}$$

$$x^2 = 0.26 * (1.79 * 10^{-5}) = 4.64 * 10^{-6}$$

$$x = 2.15 * 10^{-3}$$

$$K_b \text{ for NH}_3 = \frac{10^{-14}}{K_a} = \frac{10^{-14}}{5.6 * 10^{-10}} = 1.79 * 10^{-5}$$

$$\text{Check: } \frac{x}{0.23} = \frac{2.15 * 10^{-3}}{0.26} = 0.0083$$

$$K_b = \frac{[\text{NH}_4^+][\text{OH}^-]}{[\text{NH}_3]} = 1.79 * 10^{-5} = \frac{x * x}{0.26 - x}$$

< 0.05 so ok

Assume 0.26 >> x so 0.26 - x = 0.26

Therefore, [OH⁻] = 2.15 * 10⁻³ M

$$\text{Therefore, eqn. reduces to } 1.79 * 10^{-5} = \frac{x * x}{0.26}$$

$$\text{pOH} = -\log[\text{OH}^-] = -\log(2.15 * 10^{-3}) = 2.667$$

$$\text{pH} = 14 - \text{pOH} = 14 - 2.667 = 11.33 \text{ rounded}$$

Agree with prediction: pH > 7 as predicted

A. What is the pOH of a 0.53 M solution of ethylamine, CH₃CH₂NH₂?

•Formula:

•Type cmpd:

•Strength:

Prediction of pOH:

Reason for prediction:

Agree with prediction:

B. What is the pH of a 0.34 M solution of methylamine, CH₃NH₂?

•Formula:

•Type cmpd:

•Strength:

Prediction of pH:

Reason for prediction:

Agree with prediction:

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✓✓✓■ *Calculation of the pH or pOH of a Solution of a Weak Acid or Base AGAIN (cont'd)*

2b) C. What is the pOH of a 0.53 M solution of dimethylamine, (CH₃)₂NH?

•Formula:

•Type compd:

•Strength:

Prediction of pOH:

Reason for prediction:

Agree with prediction:

D. What is the pH of a 0.34 M solution of ammonia?

•Formula:

•Type compd:

•Strength:

Prediction of pH:

Reason for prediction:

Agree with prediction:

E. What is the pH of a 0.37 M solution of ammonia?

•Formula:

•Type compd:

•Strength:

Prediction of pH:

Reason for prediction:

Agree with prediction:

■ **An Overview of Buffers**

3a) Define buffer solution. (p. 716)

- b) A solution that contains approximately one mole of acetic acid and one mole of acetate ion can absorb either H^+ or OH^- (i.e. act like a buffer). Use LeChatelier's Principle to explain why in the chart below using arrows as in similar charts in Module 13.

<i>Stress</i> ↓	<i>Shift</i> ↓	$CH_3COOH(aq)$	+	H_2O	\rightleftharpoons	$CH_3COO^-(aq)$	+	$H_3O^+(aq)$
Add H_3O^+								
Add OH^-								

- c) What must be present for a solution to function as a buffer? (Approximately equal amounts of a weak acid and its conjugate base. Note: For our purposes, "approximately equal" means that the concentration of the one of the substances is not more than 10 times greater than the other.)

- d) Briefly describe four ways to make a buffer solution. (1: Add approximately equal amounts of a weak acid and its salt to water. 2: Add enough strong base to a weak acid to neutralize about half of the acid. 3: Add approximately equal amounts of a weak base and its salt to water. 4: Add enough strong acid to a weak base to neutralize about half of the base)

-1)

-2)

-3)

-4)

- e) The following is an initial/change/final chart for the reaction of 0.20 moles of NaOH with 0.40 moles of acetic acid. Explain why the resulting solution would like a buffer. Note: This is an example of the 2nd way to make a buffer above.

	$NaOH$	CH_3COOH	\rightarrow	$NaCH_3COO$	$+ H_2O$
initial	0.20	0.40		0	
	-0.20	-0.20		+ 0.20	
final	0	0.20		0.20	

- f) The following is an initial/change/final chart for the reaction of 0.20 moles of HCl with 0.40 moles of ammonia. Explain why the resulting solution would like a buffer. Note: This is an example of the 4th way to make a buffer above.

	HCl	NH_3	\rightarrow	NH_4Cl	$+ H_2O$
initial	0.20	0.40		0	
	-0.20	-0.20		+ 0.20	
final	0	0.20		0.20	

■ *An Overview of Buffers (continued)*

3g) Construct initial/ /final charts indicating the reaction which would occur if the following were mixed.

•Solution prepared adding 0.20 mole of acetic acid with 0.10 moles of sodium hydroxide.	•Solution prepared adding 0.20 mole of acetic acid with 0.20 moles of sodium hydroxide.

Which of the two solutions above would act like a buffer and why?

h) Why won't a solution of a strong acid and one of its salts have buffer properties?

i) Indicate whether or not the given solution would have buffer properties. Briefly explain your answer.

S. A solution which is 0.2 M in acetic acid and 0.3 M in sodium acetate.

	substance 1	substance 2	
•Formula:	CH ₃ COOH	NaCH ₃ COO	There are approximately equal concentrations of a weak acid and its salt. Therefore, the solution will act as a buffer.
•Type compd:	acid	salt	
•Strength:	weak	strong	

A. A solution which is 0.1 M in hydrofluoric acid and 0.2 sodium fluoride.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

B. A solution which is 0.2 M in hydrochloric acid and 0.2 M in sodium chloride.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

■ *An Overview of Buffers (continued)*

3i) C. A solution which is 0.3 M in aqueous ammonia and 0.4 M in ammonium chloride.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

D. A solution which is 0.2 M in hydrocyanic acid and 0.3 M sodium cyanide.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

E. A solution prepared by adding 0.5 moles of ammonia and 0.25 moles of hydrochloric acid.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

F. A solution prepared by adding 0.25 moles of acetic acid and 0.12 moles of sodium hydroxide.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

G. A solution prepared by adding 0.5 moles of nitric acid and 0.25 moles of ammonia.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

H. A solution prepared by adding 0.4 moles of hydrofluoric acid and 0.4 moles of potassium hydroxide.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

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■An Overview of Buffers (continued)

3j) Define buffer capacity. (p. 726)

k) In each of the following pairs, indicate which solution has the higher buffer capacity.

S. 50 mL of a solution which is 0.20 M in acetic acid and 0.20 M in sodium acetate or 20 mL of one which is 0.40 M in acetic acid and 0.40 M in sodium acetate?

In solution 1, $\text{mol CH}_3\text{COOH} = \text{mol NaCH}_3\text{COO} = M \cdot L = (0.20 \text{ M})(0.050 \text{ L}) = .01 \text{ mol}$

In solution 2, $\text{mol CH}_3\text{COOH} = \text{mol NaCH}_3\text{COO} = M \cdot L = (0.40 \text{ M})(0.020 \text{ L}) = .008 \text{ mol}$

Solution 1 has the greater buffering capacity since it has the larger amounts of acid/salt.

A. 70 mL of a solution which is 0.50 M in benzoic acid and 0.50 M in sodium benzoate or 60 mL of one which is 0.30 M in benzoic acid and 0.30 M in sodium benzoate?

B. 50 mL of a solution which is 0.50 M in acetic acid and 0.50 M in sodium acetate or 30 mL of one which is 0.40 M in acetic acid and 0.40 M in sodium acetate?

C. 100 mL of a solution which is 0.40 M in HF and 0.40 M in NaF or 50 mL of one which is 0.50 M in HF and 0.50 M in NaF?

D. 150 mL of a solution which is 0.40 M in HF and 0.40 M in KF or 100 mL of one which is 0.10 M in HF and 0.10 M in KF?

E. 75 mL of a solution which is 0.30 M in acetic acid and 0.30 M in potassium acetate or 100 mL of one which is 0.25 M in acetic acid and 0.25 M in potassium acetate?

✓✓✓■ Calculation of the pH of a Buffer Solution

4. Determine the pH of the solution indicated using INITIAL/CHANGE/FINAL charts as needed.

S. A solution which 0.30 M in aqueous ammonia and 0.20 M in ammonium chloride.

•Formula: $\text{NH}_3, \text{NH}_4\text{Cl}$

•Type compd: base, salt

•Strength: base weak, salt strong

Do i/ /f chart for total dissociation of NH_4Cl .Assume $0.20 \gg x$ so $0.20 + x \approx 0.20$ AND $0.30 - x \approx 0.30$. Therefore, get

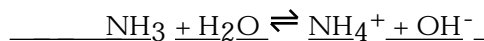
initial	0.20	0	0
	-.20	+.20	+.20
final	0	+.20	+.20

$$1.79 \times 10^{-5} = \frac{0.20 \cdot x}{0.30}$$

$$x = 0.30 \cdot (1.79 \times 10^{-5}) / 0.20$$

Do i/ /f for dissociation of NH_3 .

$$x = 2.68 \times 10^{-5}$$



initial	0.30	0.20	0
	-x	+x	+x
final	$0.30 - x$	$0.20 + x$	+x

$$\text{Check: } \frac{x}{0.20} = \frac{2.68 \times 10^{-5}}{0.20} = 0.000134$$

< 0.05 so ok

Therefore, $[\text{OH}^-] = 2.68 \times 10^{-5} \text{ M}$

$$K_b \text{ for } \text{NH}_3 = \frac{10^{-14}}{K_a} = \frac{10^{-14}}{5.6 \times 10^{-10}} = 1.79 \times 10^{-5}$$

$$\text{pOH} = -\log[\text{OH}^-] = -\log(2.68 \times 10^{-5}) = 4.572$$

$$K_b = \frac{[\text{NH}_4^+][\text{OH}^-]}{[\text{NH}_3]} = 1.79 \times 10^{-5} = \frac{(0.20 + x) \cdot x}{0.30 - x}$$

$$\text{pH} = 14 - \text{pOH} = 14 - 4.572 = 9.43 \text{ rounded}$$

*After the salt dissociates the solution contains two substances which react somewhat with water. They are NH_3 and NH_4^+ . Since the two are related (they are a conjugate acid/base pair), we only have to consider one of them (the other is automatically taken into account). Thus, in this problem, you could take into account the "weak" nature of $\text{NH}_3/\text{NH}_4^+$ by either

a) considering the dissociation which occurs when NH_3 is placed in water, i.e. $\text{NH}_3 + \text{H}_2\text{O} = \text{NH}_4^+ + \text{OH}^-$ or b) the hydrolysis of the NH_4^+ from the salt, i.e. $\text{NH}_4^+ + \text{H}_2\text{O} = \text{NH}_3 + \text{H}_3\text{O}^+$.

A. A solution which is 0.15 M in acetic acid and 0.10 M in sodium acetate.

•Formula:

•Type compd:

•Strength:

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✓✓✓■ Calculation of the pH of a Buffer Solution (continued)

4. Determine the pH of the solution indicated using INITIAL/CHANGE/FINAL charts as needed.

B. A solution which is 0.25 M in hydrofluoric acid and 0.15 M in sodium fluoride.

•Formula:

•Type compd:

•Strength:

C. A solution which is 0.40 M in aqueous ammonia and 0.10 M in ammonium chloride.

•Formula:

•Type compd:

•Strength:

✓✓✓■ *Calculation of the pH of a Buffer Solution (continued)*

4. Determine the pH of the solution indicated using INITIAL/CHANGE/FINAL charts as needed.

D. A solution which is 0.30 M in benzoic acid and 0.20 M in sodium benzoate.

•Formula:

•Type compd:

•Strength:

.....

E. A solution which is 0.25 M in propionic acid and 0.87 M in sodium propionate.

•Formula:

•Type compd:

•Strength:

.....

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✓✓✓■ Calculation of the pH of a Buffer Solution (continued)

4. Determine the pH of the solution indicated using INITIAL/CHANGE/FINAL charts as needed.

F. A solution which is 0.67 M in butylamine and 0.93 M in butylammonium chloride.

•Formula:

•Type cmpd:

•Strength:

.....

.....

■A Review of Acid-Base Titration

5a) Define titration. (pp. 161-64)

.....

b) Define equivalence point. (pp. 161, 732)

.....

c) Define indicator. (pp. 161, 745)

.....

d) Define endpoint. (The point where the indicator changes color during a titration.)

.....

e) The most common indicator is litmus. What color does litmus turn...

•in acidic solution?

•in basic solution?

.....

■ **An Review of Acid-Base Titration**

- 5f) Use the following table of common acid-base indicators to determine the color of the given indicator in the specified solution. Explain how you got your answer. (p. 749)

Indicator	pH interval for color change	Acid Color	Base Color
Methyl Violet	0-2	Yellow	Violet
Methyl Yellow	1.2-2.3	Red	Yellow
Methyl Orange	2.9-4.0	Red	Yellow
Methyl Red	4.2-6.3	Red	Yellow
Bromthymol Blue	6.0-7.6	Yellow	Blue
Thymol Blue	8.0-9.6	Yellow	Blue
Phenolphthalein	8.3-10	Colorless	Pink
Alizarin Yellow G	10.1-12.0	Yellow	Red

- S. What is the color of bromothymol blue in a solution with a pH of 11?
The pH interval over which bromothymol blue changes color is 6.0-7.6. Therefore, the indicator would be blue at pH 11 since it is above the point where is totally changed to the base color (7.6).

A. What is the color of phenolphthalein in a solution with a pH of 2?

B. What is the color of thymol blue in a solution with a pH of 10?

C. What is the color of methyl orange in a solution with a pH of 6?

D. What is the color of bromothymol blue in a solution with a pH of 8?

E. What is the color of alizarin yellow G in a solution with a pH of 4?

F. What is the color of methyl red in a solution with a pH of 7?

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■An Review of Acid-Base Titration (continued)

5g) Recommend an indicator from the list on the previous page to detect the equivalence point of the following reactions and explain your answer.

S. A titration with an equivalence point at a pH of 4.5.
Methyl red would be a good one for this titration since the equivalence point of the reaction 4.5 lies within the pH interval of change. Methyl red is red when $\text{pH} < 4.2$ and yellow when $\text{pH} > 6.3$.

A. A titration with an equivalence point at a pH of 9.2

B. A titration with an equivalence point at a pH of 10.2.

C. A titration with an equivalence point at a pH of 1.7.

D. A titration with an equivalence point at a pH of 7.3.

E. A titration with an equivalence point at a pH of 5.6.

F. A titration with an equivalence point at a pH of 3.1.

■Titration Curves

6a) Define titration curve. (736)

b) Briefly explain how buffering can affect the shape of a titration curve. (p. 739)

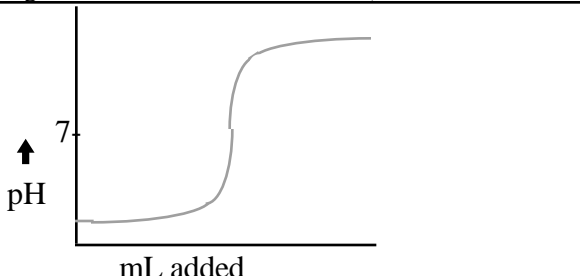
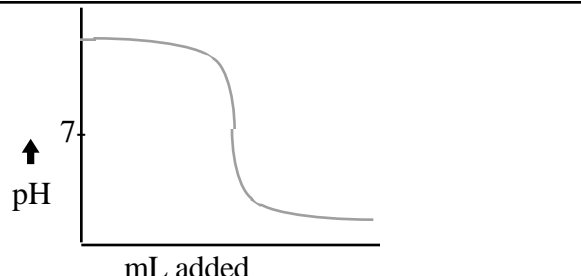
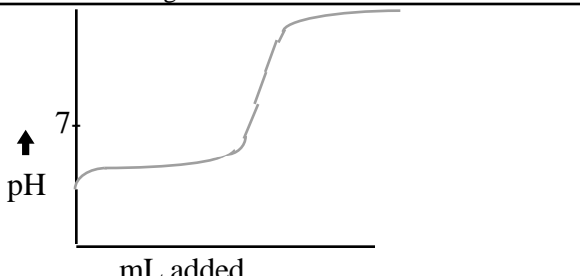
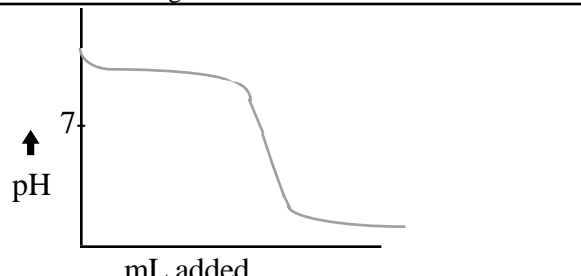
c) If a strong acid reacts with a strong base, at what pH would you expect the equivalence point to occur?

d) If a strong acid reacts with a weak base, at what pH would you expect the equivalence point to occur? Use chemical equations to explain your answer.

e) If a strong base reacts with a weak acid, at what pH would you expect the equivalence point to occur? Use chemical equations to explain your answer.

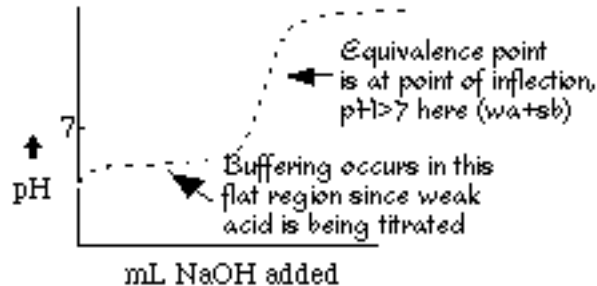
■ **Titration Curves (continued)**

6f) Below are four general types of titration curves. Identify the type of substance being titrated, the type of substance doing the titrating, the equivalence point, and any areas where buffering is occurring in each. (See Figures 15.1, 15.2, 15.3, and 15.5)

 <p>Substance in beaker: Substance being added: Substance being titrated: Substance doing titration:</p>	 <p>Substance in beaker: Substance being added: Substance being titrated: Substance doing titration:</p>
 <p>Substance in beaker: Substance being added: Substance being titrated: Substance doing titration:</p>	 <p>Substance in beaker: Substance being added: Substance being titrated: Substance doing titration:</p>

g) Sketch the titration curve for the indicated titration using the samples in 6f above. Label your axes, the equivalence point, and any region in which buffering is occurring. (Figures 15.1, 15.2, 15.3, 15.3, and 15.5)
S. Titration of 0.2 M acetic acid with sodium hydroxide.

	substance 1	substance 2	
•Formula:	CH ₃ COOH	NaOH	
•Type compd:	acid	base	
•Strength:	weak	strong	



A. Titration of 0.1 M hydrochloric acid by sodium hydroxide.

	substance 1	substance 2	
•Formula:			
•Type compd:			
•Strength:			

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■ Titration Curves (continued)

6g) Sketch the titration curve for the indicated titration. Label your axes, the equivalence point, and any region in which buffering is occurring. (Figures 15.1, 15.2, 15.3, and 15.5)

B. Titration of 0.1 M acetic acid by potassium hydroxide.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

C. Titration of 0.1 M sodium hydroxide by nitric acid.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

D. Titration of 0.1 M ammonia by hydrochloric acid.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

E. Titration of 0.1 M ammonia by nitric acid.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

F. Titration of potassium hydroxide by 0.1 M perchloric acid.

	substance 1	substance 2
•Formula:		
•Type compd:		
•Strength:		

✓✓✓■ **Calculation of the pH or pOH at a Point During a Titration—(Partial Neutralization AGAIN)**

7. F. What is the pH of a solution prepared by adding 14 milliliters of 0.42 M hydrochloric acid to 25 milliliters of 0.32 M barium hydroxide? You will need two INITIAL/CHANGE/FINAL charts.

•Formula:

•Type cmpd:

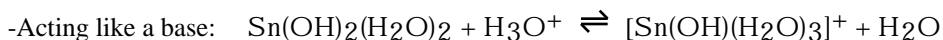
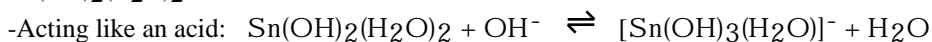
•Strength:

.....

✓✓✓■ **Amphoterism**

- 8a) Define amphoteric. (p. 660)

- b) Write a chemical equation showing the following substance acting like an acid (i.e reacting with OH⁻) and like a base (i.e reacting with H⁺).



-Acting like an acid:

-Acting like a base:



-Acting like an acid:

-Acting like a base:



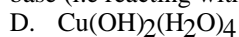
-Acting like an acid:

-Acting like a base:

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✓✓✓■Amphoterism (continued)

8b) Write a chemical equation showing the following substance acting like an acid (i.e reacting with OH^-) and like a base (i.e reacting with H^+).



-Acting like an acid:

-Acting like a base:



-Acting like an acid:

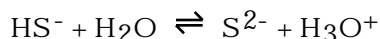
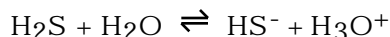
-Acting like a base:

✓✓✓■Polyprotic Acids

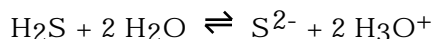
9a) Define polyprotic acid. (p. 682)

b) Write reactions showing the step-wise dissociation of the given polyprotic acid, the equilibrium expressions for each step, the overall dissociation equation, the equilibrium expression for the overall reaction, and the relationship between the step-wise and the overall dissociation constants.

S. Substance: H_2S 1) Step-wise dissociation reactions



3) Overall dissociation reaction:



5) Relationship between the answers in 2) and 4):

$$K_{\text{overall}} = K_{a1} * K_{a2}$$

2) Step-wise equil. expressions

$$K_{a1} = \frac{[\text{HS}^-][\text{H}_3\text{O}^+]}{[\text{H}_2\text{S}]}$$

$$K_{a2} = \frac{[\text{S}^{2-}][\text{H}_3\text{O}^+]}{[\text{HS}^-]}$$

4) Equilibrium expression for overall reaction

$$K_{\text{overall}} = \frac{[\text{S}^{2-}][\text{H}_3\text{O}^+]^2}{[\text{H}_2\text{S}]}$$

A. Substance: H_2CO_3 1) Step-wise dissociation reactions

2) Step-wise equil. expressions

3) Overall dissociation reaction:

4) Equilibrium expression for overall reaction

5) Relationship between the answers in 2) and 4):

✓✓✓■ **Polyprotic Acids (continued)**

9b) Write reactions showing the step-wise dissociation of the given polyprotic acid, the equilibrium expressions for each step, the overall dissociation equation, the equilibrium expression for the overall reaction, and the relationship between the step-wise and the overall dissociation constants.

B. Substance: $\text{H}_2\text{C}_2\text{O}_4$ 1) Step-wise dissociation reactions 2) Step-wise equil. expressions

3) Overall dissociation reaction:

4) Equilibrium expression for overall reaction

5) Relationship between the answers in 2) and 4):

C. Substance: H_2SO_3 1) Step-wise dissociation reactions

2) Step-wise equil. expressions

3) Overall dissociation reaction:

4) Equilibrium expression for overall reaction

5) Relationship between the answers in 2) and 4):

D. Substance: H_3PO_4 1) Step-wise dissociation reactions

2) Step-wise equil. expressions

3) Overall dissociation reaction:

4) Equilibrium expression for overall reaction

5) Relationship between the answers in 2) and 4):

E. Substance: H_3PO_3 Note: One of the H's in this compound is bonded to P and does not dissociate in water.

1) Step-wise dissociation reactions

2) Step-wise equil. expressions

3) Overall dissociation reaction:

4) Equilibrium expression for overall reaction

5) Relationship between the answers in 2) and 4):

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✓✓✓■ Polyprotic Acids (continued)

9c) The table below contains two diprotic acids. List K_{a1} and K_{a2} for each.

•H₂S: $K_{a1} =$ _____ $K_{a2} =$ _____

•H₂CO₃: $K_{a1} =$ _____ $K_{a2} =$ _____

d) What assumption is usually made when one calculates the pH of a solution of a diprotic acid such as H₂S or H₂CO₃? (p. 683)

e) What would be the predominant species present in a 0.2 M solution of H₂CO₃?

✓✓✓■ An Introduction to Formation and Dissociation Expressions

10a) Define complex ion. (p. 766)

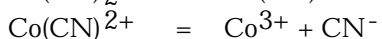
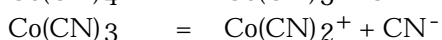
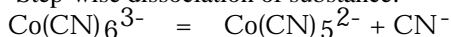
b) Define formation constant. (The equilibrium constant for the step-wise formation of a complex ion.)

c) Define dissociation constant. (The equilibrium constant for the step-wise dissociation of a complex ion.)

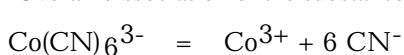
d) Write balanced step-wise reactions for either the formation or dissociation (as indicated) of the given metal ion complex. Also write equilibrium expressions for each of the equations.

S. Overall dissociation of Co(CN)₆³⁻

-Step-wise dissociation of substance:



-Overall dissociation of the substance:



-Equilibrium expressions for each step:

$$K_1 = \frac{[\text{Co(CN)}_5^{2-}][\text{CN}^-]}{[\text{Co(CN)}_6^{3-}]}$$

$$K_2 = \frac{[\text{Co(CN)}_4^-][\text{CN}^-]}{[\text{Co(CN)}_5^{2-}]}$$

$$K_3 = \frac{[\text{Co(CN)}_3][\text{CN}^-]}{[\text{Co(CN)}_4^-]}$$

$$K_4 = \frac{[\text{Co(CN)}_2^+][\text{CN}^-]}{[\text{Co(CN)}_3]}$$

$$K_5 = \frac{[\text{Co(CN)}^{2+}][\text{CN}^-]}{[\text{Co(CN)}_2^+]}$$

$$K_6 = \frac{[\text{Co}^{3+}][\text{CN}^-]}{[\text{Co(CN)}^{2+}]}$$

-Equilibrium expression for the overall reaction:

$$K_{\text{diss}} = \frac{[\text{Co}^{3+}][\text{CN}^-]^6}{[\text{Co(CN)}_6^{3-}]}$$

A. Formation of Cu(NH₃)₄²⁺

-Step-wise formation of substance:

-Equilibrium expressions for each step:

-Overall formation of the substance:

-Equilibrium expression for the overall reaction:

✓✓✓■ *An Introduction to Formation and Dissociation Expressions (continued)*

10d) Write balanced step-wise reactions for either the formation or dissociation (as indicated) of the given metal ion complex. *Also* write equilibrium expressions for each of the equations.

B. Dissociation of CuBr_4^{2-}

-Step-wise dissociation of substance:

-Equilibrium expressions for each step:

-Overall dissociation of the substance:

-Equilibrium expression for the overall reaction:

C. Formation of NiCl_4^{2-}

-Step-wise formation of substance:

-Equilibrium expressions for each step:

-Overall formation of the substance:

-Equilibrium expression for the overall reaction:

D. Dissociation of $\text{Pt}(\text{NH}_3)_4^{2+}$

-Step-wise dissociation of substance:

-Equilibrium expressions for each step:

-Overall dissociation of the substance:

-Equilibrium expression for the overall reaction:

E. Formation of HgI_4^{2-}

-Step-wise formation of substance:

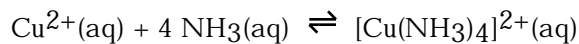
-Equilibrium expressions for each step:

-Overall formation of the substance:

-Equilibrium expression for the overall reaction:

✓✓✓■ *An Introduction to Formation and Dissociation Expressions (continued)*

10e) S. The formation constant for $[\text{Cu}(\text{NH}_3)_4]^{2+}(\text{aq})$ is 6.8×10^{12} . If you begin with similar concentrations of Cu^{2+} and NH_3 , what are the relative concentrations of $\text{Cu}^{2+}(\text{aq})$, $\text{NH}_3(\text{aq})$, and $[\text{Cu}(\text{NH}_3)_4]^{2+}(\text{aq})$ at equilibrium?



$$K_{\text{form}} = \frac{[\text{Cu}(\text{NH}_3)_4]^{2+}}{[\text{Cu}^{2+}][\text{NH}_3]^4} = 6.8 \times 10^{12}$$

$K_{\text{form}} \gg 1$, therefore $[\text{Cu}(\text{NH}_3)_4]^{2+}$ is high, $[\text{Cu}^{2+}]$ and $[\text{NH}_3]$ are low

A. The dissociation constant for $[\text{Fe}(\text{CN})_6]^{3-}$ is 1.3×10^{-37} . If you begin with similar concentrations of Fe^{3+} and CN^- , what are the relative concentrations of $\text{Fe}^{3+}(\text{aq})$, $\text{CN}^-(\text{aq})$, and $[\text{Fe}(\text{CN})_6]^{3-}(\text{aq})$ at equilibrium?

B. The formation constant for $[\text{Al}(\text{OH})_4]^-$ is 7.7×10^{33} . If you begin with similar concentrations of Al^{3+} and OH^- , what are the relative concentrations of $\text{Al}^{3+}(\text{aq})$, $\text{OH}^-(\text{aq})$, and $[\text{Al}(\text{OH})_4]^-$ at equilibrium?

C. The dissociation constant for $[\text{Cd}(\text{CN})_4]^{2-}$ is 7.7×10^{-18} . If you begin with similar concentrations of Cd^{2+} and CN^- , what are the relative concentrations of $\text{Cd}^{2+}(\text{aq})$, $\text{CN}^-(\text{aq})$, and $[\text{Cd}(\text{CN})_4]^{2-}(\text{aq})$ at equilibrium?

✓✓✓■ An Introduction to Formation and Dissociation Expressions (continued)

10e) D. The formation constant for $[\text{Ag}(\text{NH}_3)_2]^+$ is 1.6×10^7 . If you begin with similar concentrations of Ag^+ and NH_3 , what are the relative concentrations of $\text{Ag}^+(\text{aq})$, $\text{NH}_3(\text{aq})$, and $[\text{Ag}(\text{NH}_3)_2]^+(\text{aq})$ at equilibrium?

E. The dissociation constant for $[\text{HgCl}_4]^{2-}$ is 8.3×10^{-16} . If you begin with similar concentrations of Hg^{2+} and Cl^- , what are the relative concentrations of $\text{Hg}^{2+}(\text{aq})$, $\text{Cl}^-(\text{aq})$, and $[\text{HgCl}_4]^{2-}(\text{aq})$ at equilibrium?

Chemistry 1020, Module 15A

✓✓✓■ *Calculating the pH or pOH of Solutions--More practice on acid/base problems.*

11) A. What is the pH of a 0.28 M solution of acetic acid?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pH:

Reason for prediction:

Agree with prediction:

B. What is the pH of a 0.67 M solution of sodium fluoride?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pH:

Reason for prediction:

Agree with prediction:

✓✓✓■ *Calculating the pH or pOH of Solutions--More practice on acid/base problems (cont'd)*

11. C. What is the pOH of a 0.21 M solution of hydrochloric acid?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pOH:

Reason for prediction:

Agree with prediction:

D. What is the pH of a 0.49 M solution of ammonium nitrate?

•Formula:

•Type compd:

•Strength:

.....
Prediction of pH:

Reason for prediction:

Agree with prediction:

Chemistry 1020, Module 15A

✓✓✓■ *Calculating the pH or pOH of Solutions--More practice on acid/base problems (cont'd)*

11. E. What is the pH of a solution which is 0.23 M in formic acid and 0.17 M in sodium formate?

•Formula:

•Type compd:

•Strength:

.....

F. What is the pOH of a 0.37 M solution of sodium iodide?

•Formula:

•Type compd:

•Strength:

.....

Prediction of pOH:

Reason for prediction:

Agree with prediction:

.....

✓✓✓■ *Calculating the pH or pOH of Solutions--More practice on acid/base problems (cont'd)*

11. G. What is the pH of a 0.27 M solution of methylammonium chloride?

•Formula:

•Type compd:

•Strength:

Prediction of pH:

Reason for prediction:

Agree with prediction:

H. What is the pOH of a solution which is 0.37 M in ammonia and 0.29 M in ammonium chloride?

•Formula:

•Type compd:

•Strength:

Chemistry 1020, Module 15A

✓✓✓■ *Calculating the pH or pOH of Solutions--More practice on acid/base problems (cont'd)*

11. I. What is the pH of a solution which is 0.28 M in nitric acid and 0.42 M in sodium nitrate?

•Formula:

•Type compd:

•Strength:

J. What is the pH of a solution which is 0.12 M in potassium acetate and 0.32 M in acetic acid?

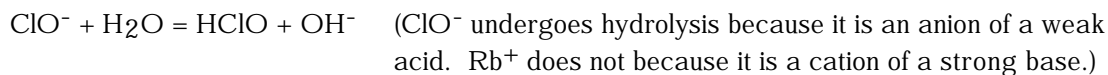
•Formula:

•Type compd:

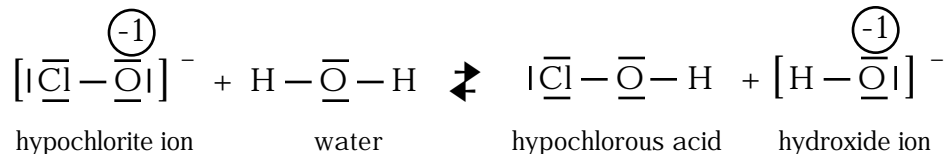
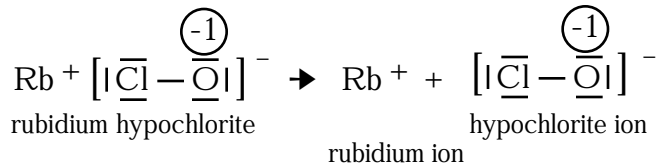
•Strength:

■ **Bonding/Model Activity to Improve Ability to Visualize in 3-D**

- S. a) Write balanced equations if rubidium hypochlorite dissociates and then undergoes hydrolysis.



- b) Draw the Lewis structure of all of the species in the above two reactions.



- c) Name all species in the equations above.
-

- A. a) Write balanced equations if sodium cyanide dissociates and then undergoes hydrolysis.

- b) Draw the Lewis structure of all of the species in the above two reactions.

- c) Name all species in the equations above.
-

- B. a) Write balanced equations if potassium acetate dissociates and then undergoes hydrolysis.

- b) Draw the Lewis structure of all of the species in the above two reactions.

- c) Name all species in the equations above.
-

Chemistry 1020, Module 15A

■Bonding/Model Activity to Improve Ability to Visualize in 3-D

C. a) Write balanced equations if ammonium chloride dissociates and then undergoes hydrolysis.

b) Draw the Lewis structure of all of the species in the above two reactions.

c) Name all species in the equations above.

D. a) Write balanced equations if potassium propionate dissociates and then undergoes hydrolysis.

b) Draw the Lewis structure of all of the species in the above two reactions.

c) Name all species in the equations above.

A-Dd) When you go to drill be prepared to assemble models of each of the reactants in the equations above and then use those models to demonstrate to your instructor what happens as the reactions occur.

■Challenge Questions

A. What is the pH of a solution prepared by adding 25 ml of 0.10 M NaOH and 50 ml of 0.10 M acetic acid?

B. What is the pH of a solution prepared by adding 15 ml of 0.10 M HCl to 50 ml of 0.10 M aqueous ammonia?

C. If the step-wise acid dissociation constants for the diprotic acid H_2Y are $K_{a1} = 2.8 \times 10^{-3}$ and $K_{a2} = 5.7 \times 10^{-8}$, what would the relative pH of a solution prepared by dissolving KHY in water. That is, would HY^- be more likely to give up a hydrogen ion or to accept one? Explain your reasoning.

D. If the step-wise acid dissociation constants for the diprotic acid H_2X are $K_{a1} = 1.4 \times 10^{-5}$ and $K_{a2} = 2.5 \times 10^{-9}$, what would the relative pH of a solution prepared by dissolving NaHX in water. That is, would HX^- be more likely to give up a hydrogen ion or to accept one? Explain your reasoning.

E. What is the percent ionization (dissociation) in a 1.00 M solution of hydrofluoric acid? Show all steps.

F. What is the percent ionization (dissociation) in a 0.86 M solution of ammonia? Show all steps.

Revised by JW Carmichael Spring, 2000; SJB 5/8/2001; SJB 5/7/2002; MA 5/5/2003