

Exam 3A

Chem 1142

Spring 2019

Name: KEY

MULTIPLE CHOICE. [2 pts ea.] Record the best response on the scantron sheet. [40 pts total.]

Assume all solutions are aqueous and at a temperature of 25 °C, unless stated otherwise.

Q1. Bases turn litmus what color?

- ☒ A) blue
- ☐ B) green
- ☐ C) red
- ☐ D) yellow

Acids turn litmus red
Bases " " blue

Q2. The Arrhenius definition of an acid is:

- ☐ A) they donate H^+ ions to other molecules
- ☒ B) they form H^+ ions in water
- ☐ C) they accept electron pairs
- ☐ D) they turn litmus red

Q3. A good example of an Arrhenius base is:

- ☐ A) NH_3
- ☒ B) $LiOH$
- ☐ C) CH_3CO_2H
- ☐ D) HCO_3^-

↳ forms OH^- in solⁿ.
 $LiOH \rightarrow Li^+ + OH^-$

Q4. The conjugate acid to HPO_4^{2-} is:

- ☐ A) H_3PO_4
- ☒ B) $H_2PO_4^-$
- ☐ C) PO_4^{3-}
- ☐ D) H_3O^+

↳ add H^+ to get conj acid

Q5. An example of a strong acid would be:

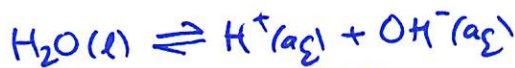
- ☐ A) HF
- ☐ B) HNO_2
- ☒ C) H_2SO_4
- ☐ D) H_3PO_4

H_2SO_4 HCl
 $HClO_4$ HBr } six strong acids!
 HNO_3 HI

Q6. At 37 °C, K_w is equal to 5.5×10^{-14} . This means the pH of pure water at this temperature will be:

- ☐ A) 7.00
- ☐ B) 5.50
- ☐ C) 13.26
- ☒ D) 6.63

$$K_w = [H^+][OH^-]$$



↑ ↑
same conc!

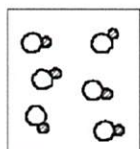
body temp... neutral
pH = 6.63!

$$-1- \Rightarrow 5.5 \times 10^{-14} = x^2 \Rightarrow x = \sqrt{5.5 \times 10^{-14}}$$

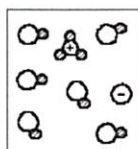
$$\Rightarrow x = 2.3 \times 10^{-7}, pH = -\log[H^+] = 6.63$$

- Q7. Which of the following diagrams represents a snapshot of a very small portion of a beaker containing a weak acid, HA, dissolved in water? Note that the solvent molecules (H_2O) are not shown for clarity.

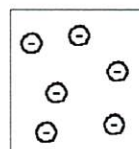
$\bigcirc\bigcirc$ = HA molecule $\bigcirc\bigcirc\bigcirc$ = H_3O^+ ion \ominus = A^- ion



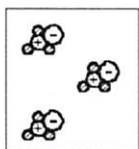
(a)



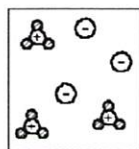
(b)



(c)

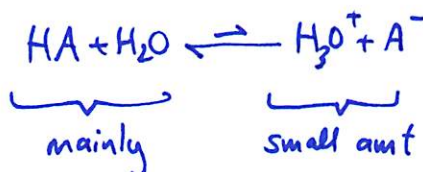


(d)



(e)

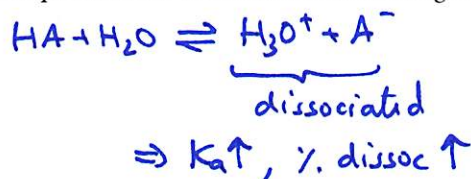
weak acids partially dissociate!



largest K_a

- Q8. Given three separate solutions containing equal concentrations of formic acid ($K_a = 1.7 \times 10^{-4}$), phenol ($K_a = 1.3 \times 10^{-10}$), and acetic acid ($K_a = 1.8 \times 10^{-5}$), select the response below that has the acids arranged in order of increasing percent dissociation at equilibrium.

- A) formic < phenol < acetic
B) formic < acetic < phenol
C) acetic < formic < phenol
D) phenol < acetic < formic



- Q9. If the concentration of hydroxide ion is $1.8 \times 10^{-4} M$ in an aqueous solution at $25^\circ C$, what is the pH of the solution?

- A) 2.74
B) 3.74
C) 9.26
D) 10.26

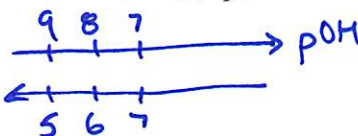
$$pOH = -\log[OH^-] = 3.74$$

$$@ 25^\circ C, pH + pOH = 14.00 \Rightarrow pH = 14.00 - pOH = 10.26$$

- Q10. Which solution has the largest pOH at $25^\circ C$: 0.100 M NaOH(aq), 0.100 M $Sr(OH)_2$ (aq), or 0.100 M HCl(aq)?

- A) 0.100 M NaOH(aq)
B) 0.100 M $Sr(OH)_2$ (aq)
C) 0.100 M HCl(aq)
D) Impossible to determine

easier way...



Big pOH = small pH = strong acid

$$pH = -\log[0.100] = 1.000$$

$$pOH = 14.00 - pH = 13.00$$

$$pOH = -\log(0.100) = 1.000$$

$$pOH = -\log(0.200) = 0.699$$



- Q11. What is the pH of 0.25 M LiOH(aq)?

- A) 0.60
B) 1.20
C) 10.25
D) 13.40



$$pOH = -\log[OH^-] = 0.60$$

$$14.00 = pH + pOH \Rightarrow pH = 13.40$$

- Q12. A Lewis base is a(n):

- A) electron-pair donor
B) proton acceptor
C) electron-pair acceptor
D) proton donor

Q13. Which pair of substances will constitute a buffer when found in solution together:

A) $\text{NaNO}_2 / \text{HNO}_3$

B) KCl / HF

C) HF / NaF

D) $\text{NaNO}_2 / \text{KNO}_2$

weak acid, HF
conj. base, F^- (Na⁺ F⁻)
counter ion

Q14. A solution containing 0.30 M $\text{HClO}(\text{aq})$ ($K_a = 3.5 \times 10^{-8}$) and 0.25 M of $\text{NaClO}(\text{aq})$ would have a pH of:

A) 7.46

B) 7.38

C) 7.54

D) 7.31

weak acid
conj. base
 ClO^-
 $\text{pK}_a = -\log(K_a) = 7.46$
H-H eq: $\text{pH} = \text{pK}_a + \log \frac{b}{a} = 7.46 + \log \frac{0.25 \text{ M}}{0.30 \text{ M}} = 7.38$

Q15. A weak monoprotic acid (HA) has a pK_a of 3.94. If we need to prepare a buffer with a pH of 3.74, then we can say for sure that:

A) $[\text{HA}] > [\text{A}^-]$

B) $[\text{HA}] = [\text{A}^-] - 0.20 \text{ M}$

C) $[\text{HA}] < [\text{A}^-]$

D) $[\text{HA}] = [\text{A}^-] + 0.20 \text{ M}$

$\text{pH} < \text{pK}_a$! So need more HA, $[\text{HA}] > [\text{A}^-]$
or: $\text{pH} = \text{pK}_a + \log \frac{b}{a} \Rightarrow \text{pH} - \text{pK}_a = -0.20 = \log \frac{b}{a}$
 $\Rightarrow \frac{[b]}{[a]} = 10^{-0.20} = 0.63$

Q16. Which of the following acids would it be best to use to prepare a buffer with a pH of 4.25?

A) HClO_2 , $K_a = 1.1 \times 10^{-2}$

B) HNO_2 , $K_a = 4.0 \times 10^{-4}$

C) HCH_2O , $K_a = 1.8 \times 10^{-4}$

D) H_2CO_3 , $K_a = 4.3 \times 10^{-7}$

$\text{pK}_a = 1.96$

$\text{pK}_a = 3.40$

$\text{pK}_a = 3.74$

$\text{pK}_a = 6.37$

$\Rightarrow [a] = \frac{1}{0.63} \times [b]$
 $(a) = 1.58 [b]$
(best buffer is when $\text{pK}_a \approx \text{pH}$)
because then $[b] \approx [a]$ $[a] > [b]$!

Q17. Which chemical equation best corresponds to the K_{sp} reaction for calcium carbonate?

A) $\text{CaCO}_3(\text{s}) \rightleftharpoons \text{Ca}^+(\text{aq}) + \text{C}^{3+}(\text{aq}) + 2\text{O}^{2-}(\text{g})$

B) $\text{CaCO}_3(\text{s}) \rightleftharpoons \text{Ca}(\text{aq}) + \text{CO}_3(\text{aq})$

C) $\text{CaCO}_3(\text{s}) \rightleftharpoons \text{Ca}^+(\text{aq}) + \text{CO}_3^{2-}(\text{aq})$

D) $\text{CaCO}_3(\text{s}) \rightleftharpoons \text{Ca}^{2+}(\text{aq}) + \text{CO}_3^{2-}(\text{aq})$

K_{sp} rxn
solid \rightleftharpoons dissolved ion

Q18. Which substance will have the **smallest** molar solubility: BaSO_4 ($K_{sp} = 1.07 \times 10^{-10}$),

CaSO_4 ($K_{sp} = 7.10 \times 10^{-5}$), or FeS ($K_{sp} = 3.72 \times 10^{-19}$).

A) BaSO_4

B) CaSO_4

C) FeS

D) Impossible to determine

$K_{sp} \downarrow$, so fewer dissolved ion
 \Rightarrow lower molar sol

Q19. In which solution would ammonium fluoride be the **most soluble**?

A) 0.25 M $\text{NH}_4\text{Br}(\text{aq})$

B) 0.10 M $\text{NH}_4\text{Cl}(\text{aq})$

C) 0.50 M $\text{LiF}(\text{aq})$

D) 0.35 M $\text{KBr}(\text{aq})$

NH_4^+ common ion

F^- common ion

no common ion! Highest sol!

$\text{NH}_4\text{F}(\text{s}) \rightleftharpoons \text{NH}_4^+(\text{aq}) + \text{F}^-(\text{aq})$

if these $[\]$'s are increased,
see shift to LHS (Le Chatelier) \Rightarrow lower sol!

Q20. The solubility product constant for magnesium hydroxide, $\text{Mg}(\text{OH})_2$ is 2.06×10^{-13} . Which solution would form a precipitate immediately upon mixing?

A) A solution containing $1.0 \times 10^{-6} \text{ M Mg}^{2+}$ and $1.0 \times 10^{-3} \text{ M OH}^-$

B) A solution containing $1.0 \times 10^{-5} \text{ M Mg}^{2+}$ and $1.0 \times 10^{-4} \text{ M OH}^-$

C) A solution containing $1.0 \times 10^{-4} \text{ M Mg}^{2+}$ and $1.0 \times 10^{-5} \text{ M OH}^-$

D) A solution containing $1.0 \times 10^{-3} \text{ M Mg}^{2+}$ and $1.0 \times 10^{-6} \text{ M OH}^-$

$\text{Mg}(\text{OH})_2(\text{s}) \rightleftharpoons \text{Mg}^{2+}(\text{aq}) + 2\text{OH}^-(\text{aq})$

$Q_{sp} = [\text{Mg}^{2+}][\text{OH}^-]^2$

if $Q_{sp} > K_{sp}$, will shift to LHS
& form ppt.

$K_{sp} \leftarrow Q_{sp}$

$Q_{sp} = (10^{-6})(10^{-3})^2 = 10^{-12} > 2.06 \times 10^{-13}$!
so will ppt upon mixing

Short Response.

Show ALL work to receive credit.

- Q21. [15 pts.] (a) The pH of 0.13 M HF(aq) is 2.01. Use this information to determine K_a for HF. Be sure to write an ICE chart as part of your answer.



I	0.13	—	≈ 0	0
C	-x	—	+x	+x
E	(0.13-x)	—	(x)	(x)

$$K_a = \frac{[\text{H}_3\text{O}^+][\text{F}^-]}{[\text{HF}]_{\text{eq}}} = \frac{x^2}{0.13-x}$$

$$\text{pH} = 2.01 \Rightarrow [\text{H}_3\text{O}^+] = 10^{-\text{pH}} = 10^{-2.01} = x \Rightarrow K_a = \frac{(10^{-2.01})^2}{0.13 - 10^{-2.01}} = 7.9 \times 10^{-4}$$

- (b) Using your calculated K_a , predict the pH of 1.3 M HF(aq).



I	1.3	—	≈ 0	0
C	-x	—	+x	+x
E	(1.3-x)	—	(x)	(x)

$$K_a = \frac{x^2}{1.3-x} = 7.9 \times 10^{-4}. \text{ Assume } x \ll 1.3$$

$$\Rightarrow 7.9 \times 10^{-4} \approx \frac{x^2}{1.3} \Rightarrow x = \sqrt{1.3 \times 7.9 \times 10^{-4}} = 0.032$$

5% rule? % ionization = $\frac{[\text{H}_3\text{O}^+]_{\text{eq}}}{[\text{HF}]_0} \times 100 = \frac{0.032}{1.3} \times 100 = 2.5\%$ (✓)

$$\begin{aligned} \text{pH} &= -\log[\text{H}^+] \\ &= -\log[0.032] \\ &= 1.49 \end{aligned}$$

- (c) What is the percent dissociation of HF in part (a) vs. part (b)?

Part (a) 7.5 %

Part (b) 2.5 %

$$\% \text{ dissociation} = \frac{x}{0.13} \times 100 = \frac{10^{-2.01}}{0.13} \times 100 = 7.5\%$$

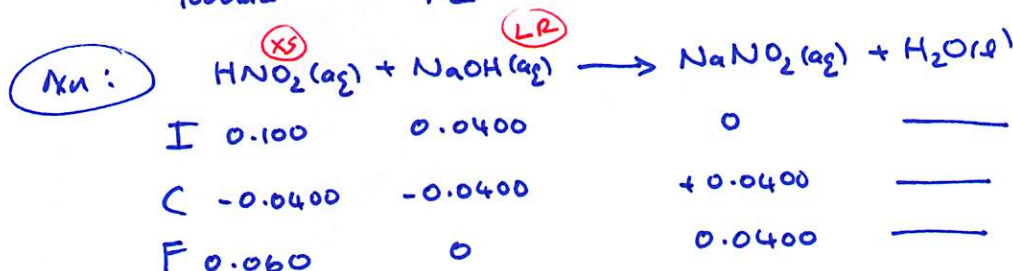
Q22. [15 pts.] (Be sure to write all relevant chemical equations and ICE/ICF charts for this question.)
 250.0 mL of 0.400 M $\text{HNO}_2(\text{aq})$, $K_a = 4.0 \times 10^{-4}$, is mixed with 50.0 mL of 0.800 M $\text{NaOH}(\text{aq})$.

(a) Calculate the pH of the solution formed.

$$\# \text{ mol HNO}_2 = 250.0 \text{ mL} \times \frac{1 \text{ L}}{1000 \text{ mL}} \times \frac{0.400 \text{ mol HNO}_2}{1 \text{ L}} = 0.100 \text{ mol HNO}_2$$

$$\text{p}K_a = -\log(K_a) = 3.40$$

$$\# \text{ mol NaOH} = 50.0 \text{ mL} \times \frac{1 \text{ L}}{1000 \text{ mL}} \times \frac{0.800 \text{ mol NaOH}}{1 \text{ L}} = 0.0400 \text{ mol NaOH}$$

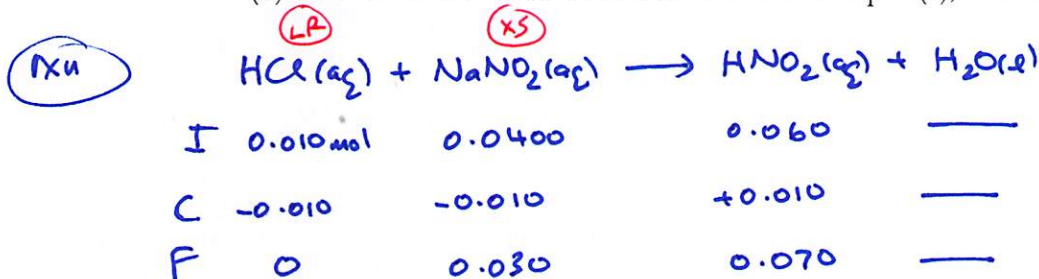


Buffer!
 (weak acid/conj. base)

$$\text{pH} = \text{p}K_a + \log \frac{[\text{base}]}{[\text{acid}]} = 3.40 + \log \frac{0.0400 \text{ mol} / 0.3000 \text{ L}}{0.060 \text{ mol} / 0.3000 \text{ L}} = \boxed{3.22}$$

cancels out!

(b) If 0.010 mol HCl is added to the solution formed in part (a), what will the final pH be?



Buffer!

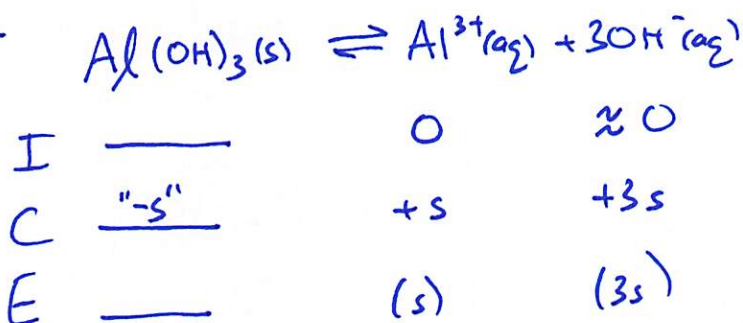
$$\text{pH} = \text{p}K_a + \log \frac{[\text{base}]}{[\text{acid}]} = 3.40 + \log \frac{0.030}{0.070} = \boxed{3.03}$$



Q23. [15 pts.] Calculate the molar solubility of aluminum hydroxide in pure water vs. in 0.200 M KOH(aq). Comment on the difference. K_{sp} for $Al(OH)_3$ is 2.3×10^{-8} .

Be sure to write the K_{sp} chemical equation and an ICE chart as part of your answer! Explain any assumptions you are making.

in water
 K_{sp} rxn:

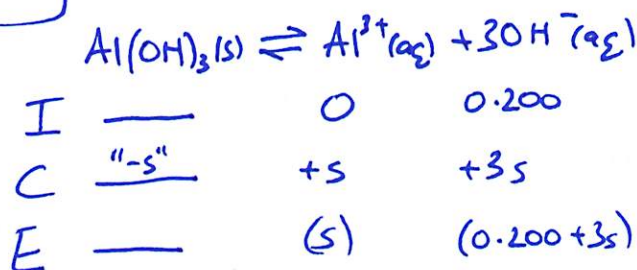
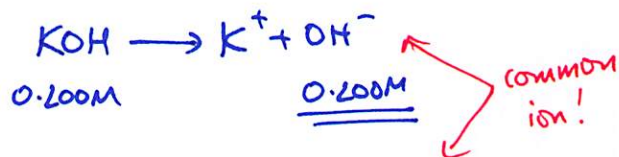


$$K_{sp} = [Al^{3+}][OH^{-}]^3$$

$$\Rightarrow 2.3 \times 10^{-8} = (s)(3s)^3 = 27s^4$$

$$\Rightarrow s = \sqrt[4]{\frac{2.3 \times 10^{-8}}{27}} = \boxed{5.4 \times 10^{-3} M}$$

when dissolving in 0.200 M KOH, there's a common ion, OH^{-} !



$$K_{sp} = [Al^{3+}][OH^{-}]^3$$

$$\Rightarrow 2.3 \times 10^{-8} = (s)(0.200 + 3s)^3 \approx (s)(0.200)^3$$

$$\Rightarrow s \approx \frac{2.3 \times 10^{-8}}{(0.200)^3} = \boxed{2.9 \times 10^{-6} M}$$

(so, yes! $3s \ll 0.200$)

THE EFFORT TO AVOID 'CHEM-SPEAK'



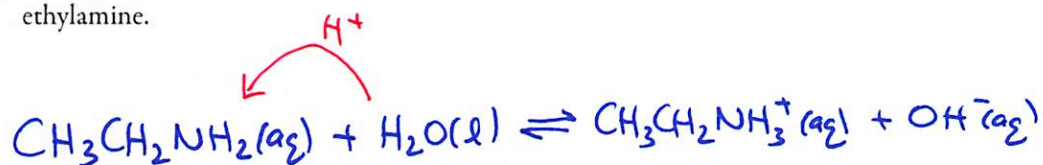
comment

molar sol. is much lower when OH^{-} is already present.

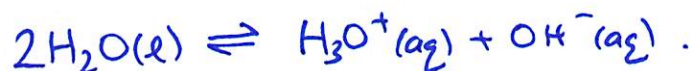
This is expected due to Le Chatelier!

Inc. $[OH^{-}]$, causes shift to LHS... so less ppt(s) will dissolve!

- Q24. [15 pts.] (a) Write out the chemical equation (reaction) corresponding to K_b for $\text{CH}_3\text{CH}_2\text{NH}_2(\text{aq})$, ethylamine.



- (b) Write out the chemical equation (reaction) corresponding to K_w .

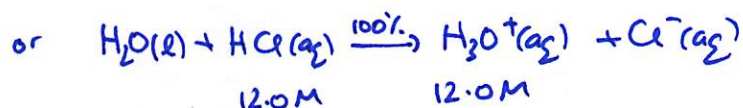


- (c) Without performing a calculation, explain how the pH of 0.100 M $\text{NH}_3(\text{aq})$ compares with 0.100 M $\text{LiOH}(\text{aq})$.

LiOH is a strong base (100% dissociation), so will have a greater pH than NH_3 , a weak base of same conc.

- (d) Calculate the pH of 12.0 M $\text{HCl}(\text{aq})$ (concentrated hydrochloric acid).

— strong acid!



$$\text{pH} = -\log [\text{H}^+] = -\log [12.0] = -1.079$$

(3sf.) (3dp)

Bonus question

Predict whether the following aqueous salts will be acidic, basic, neutral, or whether there is not enough information to decide.

NaF basic

$\text{Al}(\text{NO}_3)_3$ acidic

LiCl neutral

Na^+ : neutral
(from strong base, NaOH)

F^- : basic
(from weak acid, HF)

...

Periodic Table of the Elements

IA	IIA											IIIA	IVA	VA	VIA	VIIA	VIIIA
1 H 1.008	2 He 4.003																
3 Li 6.941	4 Be 9.012											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.31											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.07	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.87	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.61	33 As 74.92160	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc [98]	44 Ru 101.1	45 Rh 102.9	46 Pd 106.4	47 Ag 107.9	48 Cd 112.4	49 In 114.8	50 Sn 118.7	51 Sb 121.8	52 Te 127.60	53 I 126.9	54 Xe 131.3
55 Cs 132.9	56 Ba* 137.3	57 La 138.9	58 Ce 140.1	59 Pr 140.9	60 Nd 144.2	61 Pm [145]	62 Sm 150.4	63 Eu 152.0	64 Gd 157.3	65 Tb 158.9	66 Dy 162.50	67 Ho 164.9	68 Er 167.3	69 Tm 168.9	70 Yb 173.0		
87 Fr [223]	88 Ra** [226]	89 Ac [227]	90 Th 232.0	91 Pa 231.0	92 U 238.0	93 Np [237]	94 Pu [244]	95 Am [243]	96 Cm [247]	97 Bk [247]	98 Cf [251]	99 Es [252]	100 Fm [257]	101 Md [258]	102 No [259]		

$$N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$$

$$R = 8.3145 \frac{\text{J}}{\text{mol} \cdot \text{K}} = 0.08206 \frac{\text{atm} \cdot \text{L}}{\text{mol} \cdot \text{K}}$$

$$M_1 V_1 = M_2 V_2$$

$$\text{pH} = -\log_{10}[\text{H}^+]$$

$$\text{pH} + \text{pOH} = 14.00 \text{ (25 } ^\circ\text{C)}$$

$$K_w = 1.0 \times 10^{-14} \text{ (25 } ^\circ\text{C)}$$

$$K_a \cdot K_b = K_w$$

$$\text{pH} = \text{p}K_a + \log \frac{[\text{Base}]}{[\text{Acid}]}$$

$$\text{Given: } ax^2 + bx + c = 0, \text{ then } x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$$

