Unit 2

PREPARATION OF SOLUTIONS (CONCENTRATIONS)

- 1- The mw of Na₂CO₃ is : Na=23, O=16, C=12
- A) 140
- B) 106
- C) 96
- D) 100
- E) 60

- 2- How many grams of Na₂CO₃ in 3 moles, (mw = 106)
- A) 318
- B) 0.028
- C) 134
- D) 201
- E) 67

- 3- Calculate the normal concentration (N) of 0.1 M solution of HCl
- A) 0.2 N
- B) 0.3 N
- C) 0.05 N
- D) 0.1 N

- 4- The eq.wt of H_3PO_4 (mw=98) is:
- A) 98
- B) 49
- C) 32.7
- D) 196
- E) 294

 5- 2.5 g of Na₂SO₄ (mw = 106) has been dissolved in water and the volume was completed to 500 mL, calculate the followings :

The molar concentration of Na₂SO₄

A) 5

- B) 500
- C) 0.5
- D) 0.05

The limiting reactant is:

- A) present in lower of moles
- B) present in higherof moles
- C) present in lower of mass
- D) present in lower of mass

The limiting reactant is:

A) It is comletly used up in the reaction

B) It is not comletly used up in the reaction

In chemical equilibrium:

- A) rate (forward= reverse)
- B) rate (forward< reverse)
- C) rate (forward> reverse)

If Keq >> 1 (M=0.1 of product)

- A) Product 0.1-x
- B) product 0.1+x

If Keq << 1 (M=0.1 of product)

- A) Product 0.1-x
- B) Product x

Calculate the molar concentration of [H⁺] in 0.16 M solution of dichloroacetic acid Cl₂CHCOOH

- ($Ka = 5 \times 10^{-3}$). The concentration of the acid is initial (before dissociation)
- $Cl_2CHCOOH \rightarrow Cl_2CHCOO + H^+$
- The concentration of Cl_2CHCOO and H^+
- A) 0.16-x
- B) 0.16+x
- C) X

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- $Cl_2CHCOOH \rightarrow Cl_2CHCOO- + H^+$
- The concentration of Cl₂CHCOOH and H⁺
- A) 0.16-x
- B) 0.16+x
- C) X

Calculate the molar concentration of [H⁺] in 0.16 M solution of dichloroacetic acid Cl₂CHCOOH

- ($Ka = 5 \times 10^{-3}$). The concentration of the acid is initial (before dissociation)
- $Cl_2CHCOOH \rightarrow Cl_2CHCOO- + H^+$
- The value of x
- A) 5x10⁻³
- B) 0.16
- C) 5.9x10³

- Acid and base strong
- A) high conc.
- B) high vol.
- C) Completely dissociate

- рΗ
- A) 1-7
- B) 1-14
- C) 7-14
- D) 0-14

- POH
- A) 1-7
- B) 1-14
- C) 7-14
- D) 0-14

pH of HCl(0.01M) A)pH= 1 B) pH=2 pH= 3

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pOH of Ba(OH)2 (0.01M)
A)pOH= 0.7
B) pOH=1.7
C) pOH= 3
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pOH of NaOH(0.03M)
A)pOH= 1
B) pOH=1.5
pOH= 3
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pOH of HCl(0.01M) A)pOH= 2 B) pOH=5 pOH= 12

pH of strong acid :

- A) $pH = -\log[H^+]$
- B) $pH = -log\sqrt{K_a C_a}$
- C) $pH = -log\sqrt{K_b C_b}$

- pH of strong base :
- A) $pH = -\log[H^+]$
- B) $pOH = -\log[OH^{-}]$
- C) $pH = -log\sqrt{K_a C_a}$

- pH of weak acide :
- A) $pH = -\log[H^+]$
- B) $pOH = -\log[OH^{-}]$
- C) $pH = -log\sqrt{K_a C_a}$

pH of weak base:

- A) $pH = -\log[H^+]$
- B) $pH = -log\sqrt{K_a C_a}$
- C) $pH = -log\sqrt{K_b C_b}$

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

B) $K_w = [H^+] [OH^-]$
C) $K_w = [H^+] - [OH^-]$
D) $K_w = \frac{[H^+]}{[OH^-]}$

Κ

- D) $pK_w = pH pOH$
- C) $pK_w = pH + pOH$
- B) $pK_w = [H^+] + [OH^-]$
- A) $pK_w = [H^+][OH^-]$
- pK_w

Calculate the pH of. 0.5 M solution of NH_4Cl ? Kb (NH_3) = 1.75 X 10⁻⁵ A) $pH = -\log[H^+]$

- **B)** $pH = -log\sqrt{K_a C_a}$
- $\mathbf{C} \quad pH = -\log \sqrt{\frac{K_w C_s}{K_b}}$
- D) $pH = -log\sqrt{K_{la} C_s}$

Calculate the pH of. 0.5 M solution of NH₄Cl ? Kb (NH₃) = 1.75 X 10⁻⁵ A) $pH = -\log [H^+]$ B) $pH = -\log \sqrt{K_a C_a}$ C) $pH = -\log \sqrt{\frac{K_w C_s}{K_b}}$

D) $pH = -log\sqrt{K_{la} C_s}$

Calculate the pH of. 0.5 M solution of NH_4Cl ? Kb (NH_3) = 1.75 X 10⁻⁵

A) pH = 2.1
B) pH = 4.8
C) pH = 7
D) pH = 9.6

Calculate the pH of 0.2 M solution of Ba(OH)²

A) pOH = 12.1
B) pOH = 8.9
C) pOH = 5.5
D) pOH = 0.4

- Calculate the pH of 0.2 M solution of Ba(OH)2 •
- A) 9.6
- B) 11.6
- C) 13.6

pH of salt (strong acid+ weak base)

- A) pH=7
- B) $pH = -log\sqrt{K_a C_a}$
- $\mathsf{C}) \quad \overset{pH}{=} \quad -\log\left[H^+\right]$
- $\mathsf{D}) \quad pH = -\log \sqrt{\frac{K_w C_s}{K_b}}$

pH of salt (strong acid+ strong base) A) pH=7 B) $mH = -log \frac{K_w C_s}{K_w C_s}$

B) $pH = -log \sqrt{\frac{K_w C_s}{K_b}}$ C) $pOH = -log \sqrt{\frac{k_w c_s}{k_a}}$

pH of salt (strong base + weak acid) A) pH=7 B) $pH = -log \sqrt{\frac{K_w C_s}{K_b}}$ C) $pOH = -log \sqrt{\frac{k_w C_s}{k_a}}$

pH of CH₃COOH /CH₃COONa
A)
$$pOH = pK_b + log \frac{C_s}{C_b}$$

B) $pH = pK_a + log \frac{C_s}{C_a}$
C) $pOH = -log \sqrt{\frac{k_w C_s}{k_a}}$

pH of NH₄OH / NH₄Cl
A)
$$pOH = pK_b + log \frac{C_s}{C_b}$$

B) $pH = pK_a + log \frac{C_s}{C_a}$
C) $pOH = -log \sqrt{\frac{k_w C_s}{k_a}}$

Calculate the pH of a solution containing 0.1 M Na_2CO_3 and 0.2 M $NaHCO_3$? Ka2 (H_2CO_3) = 4.7 X10⁻¹¹.

- A) pH= 5
- B) pH= 8
- C) pH=10

Calculate the pH of a solution resulting from adding 4 mL of 0.2 M of NaOH solution to 16 mL of a buffer solution containing 0.1 M CH₃COOH

- ($K_a = 1.8X10^{-5}$) and 0.2 M CH₃COONa
- The number of mmol CH_3COOH
- A) 0.8 mmol
- B) 1.6 mmol
- C) 4 mmol
Unite 5: BUFFER SOLUTIONS

Calculate the pH of a solution resulting from adding 4 mL of 0.2 M of NaOH solution to 16 mL of a buffer solution containing 0.1 M CH₃COOH

($K_a = 1.8X10^{-5}$) and 0.2 M CH₃COONa

The number of mmol NaOH

A) 0.8 mmol

- B) 1.6 mmol
- C) 4 mmol

Unite 5: BUFFER SOLUTIONS

Calculate the pH of a solution resulting from adding 4 mL of 0.2 M of NaOH solution to 16 mL of a buffer solution containing 0.1 M CH₃COOH

- ($K_a = 1.8X10^{-5}$) and 0.2 M CH₃COONa
- The number of mmol CH₃COONa
- A) 0.8 mmol
- B) 1.6 mmol
- C) 4 mmol

Unite 5: BUFFER SOLUTIONS

Calculate the pH of a solution resulting from adding 4 mL of 0.2 M of NaOH solution to 16 mL of a buffer solution containing 0.1 M CH₃COOH

($K_a = 1.8X10^{-5}$) and 0.2 M CH₃COONa

The pH of a solution

- A) pH = 2
- B) pH= 5.4
- C) pH= 9.4

In the case of titrating 10 mL solution of 0.1 M of Cl⁻ (in the conical flask) by 0.2 M solution of Ag⁺ (in the burette) , calculate Veq.p of Ag⁺ solution

- A)2.5 mL
- B)5 mL
- C)10 mL

In the case of titrating 10 mL of 0.1 M of H_2SO_4 solution by 0.2 M of NaOH solution , calculate Veq.p of NaOH solution

A)2.5 mL

B)5 mL

C)10 mL

Endpoint is

- A) The same (equal) equivalence point
- B) Before equivalence point
- C) After equivalence point

300 mg of a Na₂C₂O₄ (mw = 134) 95 %w/w pure reagent was transferred to a titration conical flask . After adding acid solution and a suitable indicator , $C_2O_4^{2-}$ was titrated with KMnO4 unknown solution according to the following titration reaction equation :

2 KMnO₄ + 5 C₂O₄ + 16H+ ----- 2Mn2+ + 10CO2 + 8H2O

If the volume of KMnO₄ solution at the equivalent point was 34 mL, calculate the molarity of KMnO₄ solution, <u>the pure wieght of Na₂C₂O₄</u>

A) 300 mg

B) 285 mg

C) 31.67 mg

300 mg of a Na₂C₂O₄ (mw = 134) 95 %w/w pure reagent was transferred to a titration conical flask . After adding acid solution and a suitable indicator , $C_2O_4^{2-}$ was titrated with KMnO4 unknown solution according to the following titration reaction equation :

2 KMnO₄ + 5 C₂O₄ + 16H+ ----- 2Mn2+ + 10CO2 + 8H2O

If the volume of $KMnO_4$ solution at the equivalent point was 34 mL, calculate the molarity of $KMnO_4$ solution, the moles of $Na_2C_2O_4$.

A) 5.61 mol

B) 2.12 mol

C) 0.85 mol

300 mg of a Na₂C₂O₄ (mw = 134) 95 %w/w pure reagent was transferred to a titration conical flask . After adding acid solution and a suitable indicator , $C_2O_4^{2-}$ was titrated with KMnO4 unknown solution according to the following titration reaction equation :

2 KMnO₄ + 5 C₂O₄ + 16H+ ----- 2Mn2+ + 10CO2 + 8H2O

If the volume of $KMnO_4$ solution at the equivalent point was 34 mL, calculate the molarity of $KMnO_4$ solution, <u>the moles of $KMnO_4$ </u>

A) 5.61 mol

B) 2.12 mol

C) 0.85 mol

750 mg of a sample containing iron (aw = 55.85) was transferred into a titration conical flask and all iron was converted to Fe³⁺, then unknown excess of KI was added and the following reaction was occurred

 $2 I^{-} + 2 Fe^{3+} \rightarrow I2 + 2 Fe^{2+}$

The iodine I2 formed was titrated with 0.075 M of Na2S2O3 solution using starch as indicator according to the following reaction equation :

$$S_4O_6^{2-} + 2I_2 \rightarrow S_2O_3^{2-} + 2I_2$$

mmol of S₄O₆²⁻

A) 18.51

B) 1.39

C) 4.77

750 mg of a sample containing iron (aw = 55.85) was transferred into a titration conical flask and all iron was converted to Fe³⁺, then unknown excess of KI was added and the following reaction was occurred

 $2 |^{-} + 2 \operatorname{Fe}^{3+} \rightarrow |2 + 2 \operatorname{Fe}^{2+}$

The iodine I2 formed was titrated with 0.075 M of Na2S2O3 solution using starch as indicator according to the following reaction equation :

$$S_4O_6^{2-}+2I_2 \rightarrow S_2O_3^{2-}+2I^{-}$$

mmol of I₂

- A) 18.51 B) 1.39
- C) 0.69

750 mg of a sample containing iron (aw = 55.85) was transferred into a titration conical flask and all iron was converted to Fe³⁺, then unknown excess of KI was added and the following reaction was occurred

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$$S_4O_6^{2-}+2I_2 \rightarrow S_2O_3^{2-}+2I^{-}$$

mmol of Fe³⁺

- A) 18.51
- B) 1.39
- C) 0.69

1.782 g of a sample containing Cr^{6+} (aw = 52) is dissolved in a conical flask . An excess of 1.5 g of Fe²⁺ (aw = 55.85) are added which is oxidized to Fe³⁺ during its reduction of Cr⁶⁺ to Cr³⁺ . The excess Fe²⁺ is titrated with 0.05 N of K₂Cr₂O₇ and was found to require 10 mL at the equivalent point . Calculate the %w/w percentage of Cr⁶⁺ in the sample .

m.eq. of K₂Cr₂O₇

- A) 0.05 m.eq
- B) 0.5 m.eq
- C) 1.5 m.eq

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m.eq. of Fe²⁺ (excess) which reacted with K₂Cr₂O₇ A) 0.05 m.eq

- B) 0.5 m.eq
- C) 1.5 m.eq

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m.eq. of Fe²⁺ total

- A) 0.05 m.eq
- B) 0.78 m.eq
- C) 26.85 m.eq

1.782 g of a sample containing Cr^{6+} (aw = 52) is dissolved in a conical flask . An excess of 1.5 g of Fe²⁺ (aw = 55.85) are added which is oxidized to Fe³⁺ during its reduction of Cr^{6+} to Cr^{3+} . The excess Fe²⁺ is titrated with 0.05 N of $K_2Cr_2O_7$ and was found to require 10 mL at the equivalent point . Calculate the %w/w percentage of Cr^{6+} in the sample .

m.eq. of Fe²⁺ which reacted with Cr⁶⁺

- A) 0.05 m.eq
- B) 0.78 m.eq
- C) 26.36 m.eq

1.782 g of a sample containing Cr^{6+} (aw = 52) is dissolved in a conical flask . An excess of 1.5 g of Fe²⁺ (aw = 55.85) are added which is oxidized to Fe³⁺ during its reduction of Cr^{6+} to Cr^{3+} . The excess Fe²⁺ is titrated with 0.05 N of $K_2Cr_2O_7$ and was found to require 10 mL at the equivalent point . Calculate the %w/w percentage of Cr^{6+} in the sample .

m.eq. of Cr⁶⁺

- A) 0.05 m.eq
- B) 0.78 m.eq
- C) 26.36 m.eq

1.782 g of a sample containing Cr^{6+} (aw = 52) is dissolved in a conical flask . An excess of 1.5 g of Fe²⁺ (aw = 55.85) are added which is oxidized to Fe³⁺ during its reduction of Cr^{6+} to Cr^{3+} . The excess Fe²⁺ is titrated with 0.05 N of $K_2Cr_2O_7$ and was found to require 10 mL at the equivalent point . Calculate the %w/w percentage of Cr^{6+} in the sample .

mass of Cr⁶⁺

- A) 17.33 mg
- B) 456.91 mg
- C) 630.02 mg

1.782 g of a sample containing Cr^{6+} (aw = 52) is dissolved in a conical flask . An excess of 1.5 g of Fe^{2+} (aw = 55.85) are added which is oxidized to Fe^{3+} during its reduction of Cr^{6+} to Cr^{3+} . The excess Fe^{2+} is titrated with 0.05 N of $K_2Cr_2O_7$ and was found to require 10 mL at the equivalent point . Calculate the %w/w percentage of Cr^{6+} in the sample .

- A) 17.33 %
- B) 40.91 %
- C) 25.60%

10 mL of 0.1 M of analyte A were titrated with 0.2 M of titrant B according to the following titration reaction equation :

A + 2 B \leftrightarrow 3 E , Keq = 1x10¹⁰

Calculate the molar concentration of each [A] , [B] and [E] in the conical flask after the following additions of the titrant B solution : (1) 5 mL (2) 10 mL (3) 15 mL

calculate the **volume of the titrant B** solution at the equivalent point

A) 5 mL

B) 2.5mL

C) 10 mL

10 mL of 0.1 M of analyte A were titrated with 0.2 M of titrant B according to the following titration reaction equation :

A + 2 B \leftrightarrow 3 E , Keq = 1x10¹⁰

Calculate the molar concentration of each [A], [B] and [E] in the conical flask after the following additions of the titrant B solution : (1) 3 mL (2) 10 mL (3) 15 mL calculate the molar concentration of [E] A) 1 mL B) 0.07mL C) 0.1 mL

10 mL of 0.1 M of analyte A were titrated with 0.2 M of titrant B according to the following titration reaction equation :

A + 2 B \leftrightarrow 3 E , Keq = 1x10¹⁰

Calculate the molar concentration of each [A], [B] and [E] in the conical flask after the following additions of the titrant B solution : (1) 3 mL (2) 10 mL (3) 15 mL calculate the molar concentration of [A] A) 1 M B) 0.03 M C) 0.05 M

10 mL of 0.1 M of analyte A were titrated with 0.2 M of titrant B according to the following titration reaction equation :

A + 2 B \leftrightarrow 3 E , Keq = 1x10¹⁰

Calculate the molar concentration of each [A], [B] and [E] in the conical flask after the following additions of the titrant B solution : (1) 3 mL (2) 10 mL (3) 15 mL

calculate the molar concentration of [B]

- A) 0.3 M
- B) 1.8x10⁻⁶ M
- C) 8.3x10⁻⁷

If 10 mL of 0.1 M CH_3COOH ((Ka = 1.8 x 10⁻⁵) is titrated with 0.2 M NaOH solution , calculate the pH of the titration solution in the conical flask after the addition of the following volumes of NaOH solution : V.eq.p

- A) 5
- B) 10
- C) 15

If 10 mL of 0.1 M CH_3COOH ((Ka = 1.8 x 10⁻⁵) is titrated with 0.2 M NaOH solution, calculate the pH of the titration solution in the conical flask after the addition of the following volumes of NaOH solution : **3mL**

- A) 2.91
- B) 4.96
- C) 11.51

If 10 mL of 0.1 M CH_3COOH ((Ka = 1.8 x 10⁻⁵) is titrated with 0.2 M NaOH solution , calculate the pH of the titration solution in the conical flask after the addition of the following volumes of NaOH solution : **5mL**

- A) 5.2
- B) 8.8
- C) 11.51

If 10 mL of 0.1 M CH_3COOH ((Ka = 1.8 x 10⁻⁵) is titrated with 0.2 M NaOH solution , calculate the pH of the titration solution in the conical flask after the addition of the following volumes of NaOH solution : **10mL**

- A) 2.91
- B) 1.30
- C) 12.71

A) if pKa = 3, so, sharper the region of the curve near the equivalent point

B) if pKa = 7, so, sharper the region of the curve near the equivalent point

C) if pKa = 10, so, sharper the region of the curve near the equivalent point

At a low pH, the human can distinguish the acidic color if:

A) $\frac{[In]}{[HIn]} = \frac{10}{1}$ B) $\frac{[HIn]}{[In]} = \frac{10}{1}$ C) $\frac{[In]}{[HIn]} = \frac{20}{1}$

At a high pH, the human can distinguish the acidic color if:

A)
$$\frac{[In]}{[HIn]} = \frac{10}{1}$$

B)
$$\frac{[\text{HIn}]}{[\text{In}]} = \frac{10}{1}$$

C) $\frac{[In]}{[\text{HIn}]} = \frac{20}{1}$

At low pH (acidic), The pH of indicator is :

A) $pH_{ln} = pK_{Hln}-1$ B) $pH_{ln} = pK_{Hln}+1$ C) $pK_{Hln} = pH_{ln}-1$

The Indicator's Range is:

- A) $\Delta pK_{HIn} = pK_{HIn} \pm 1$
- B) $\Delta pK_{HIn} = pH_{In} \pm 1$
- C) $\Delta pH_{In} = pK_{HIn} \pm 1$

How do the following indicators behave with the titration of 10 mL of 0.05 M of the weak acid HA ($K_a = 1 \times 10^{-5}$) with 0.1 M NaOH : (1) $pK_{ln} = 7$ (2) $pK_{ln} = 9$ (3) $pK_{ln} = 11$ **Calculate V**_{eq.p.} A) 2.5 mL B) 5mL

C) 7.5mL

How do the following indicators behave with the titration of 10 mL of 0.05 M of the weak acid HA ($K_a = 1 \times 10^{-5}$) with 0.1 M NaOH : (1) $pK_{ln} = 7$ (2) $pK_{ln} = 9$ (3) $pK_{ln} = 11$ Calculate the pH at eq.p.

A) 4.7 mL

B) 9.3mL

C) 12.5mL

How do the following indicators behave with the titration of 10 mL of 0.05 M of the weak acid HA ($K_a = 1 \times 10^{-5}$) with 0.1 M NaOH : (1) $pK_{ln} = 7$ (2) $pK_{ln} = 9$ (3) $pK_{ln} = 11$ The suitable indicatore is

How do the following indicators behave with the titration of 20 mL of 0.1 M of NH₃ ($K_b \approx 2 X 10^{-5}$) with 0.4 M HCl :

- (1) $pK_{ln} = 3$ (2) $pK_{ln} = 5$ (3) $pK_{ln} = 7$ Calculate $V_{eq.p.}$
- A) 2.5 mL
- B) 5mL
- C) 7.5mL
Unite 8: ACID – BASE TITRATION INDICATORS

How do the following indicators behave with the titration of 20 mL of 0.1 M of NH_3 ($K_b \approx 2x \ 10^{-5}$) with 0.4 M HCl :

(1) $pK_{ln} = 3$ (2) $pK_{ln} = 5$ (3) $pK_{ln} = 7$

Calculate the pH at eq.p.

- A) 11.9
- B) 9.8

C) 5.2

Unite 8: ACID – BASE TITRATION INDICATORS

How do the following indicators behave with the titration of 20 mL of 0.1 M of NH₃($K_b \approx 2x \ 10^{-5}$) with 0.4 M HCl :

(1) $pK_{ln} = 3$ (2) $pK_{ln} = 5$

The suitable indicatore is

A) pK_{In} = 3 B) pK_{In} = 5 C) pK_{In} = 7

Unite 9: COMPLEX FORMATION TITRATION

Example of unidentate A) H₂NCH₂CH₂NH₂ B) Br⁻, F⁻, I⁻, CN-, RNH2, RCOO C) EDTA

Unite 9: COMPLEX FORMATION TITRATION

- The charge of complex Fe^{3+} , $Fe(CN)_6$
- A) 3+
- B) 3-
- C) 6-
- D) 6+

Unite 9: COMPLEX FORMATION TITRATION

Calculate the molar concentration of $[CN^{-}]$ in a solution of Cu(CN)₄²⁻ (Kd = 5.2 X 10⁻²⁸) prepared by dissolving 0.05 mole of the complex in water and completing the volume to one liter A) 2.52x10 10⁻⁵

B) 5.63x10⁻²

C) 32.2x10⁻²⁸