Exercise 2(Going step-by-step)

# Acid and Bases

1.	$C_2H_5OH$ ionises as						
	a) $C_2H_5OH \longleftarrow C_2H_5O^- + H^+$						
	b) $2C_2H_5OH \longleftarrow C_2H_5OH_2 + C_2H_5O^-$						
	c) $C_2H_5OH \longrightarrow C_2H_5^+ + OH^-$						
	d) $C_2H_5OH + H_2O \longleftarrow C_2H_5O^{} + H_3O^+$						
2.	$C_2H_5Ona$ is ain $C_2H_5OH$ .						
	a) <b>Strong base</b> b) Strong acid						
	c) Weak acid d) weak base						
3.	NH <sub>4</sub> NO <sub>3</sub> is a						
	a) Strong base in $NH_3$ b) weak base in $NH_3$						
	c) <b>Strong acid in NH</b> <sub>3</sub> d) weak acid in NH <sub>3</sub>						
4.	Select the correct statement(s).						
	a) Arrhenius theory is restricted to aqueous solution						
	b) Franklin's theory is applicable also to other ionisable						
	c) Both a&b						
	d) None of the above						
5.	In the following equilibrium,						
	$B + H_2O \iff BH^+ + OH^-$						
	a) Arrhenius acid-base concept is observed						
	b) Franklins acid –base concept is observed						
	c) Both a&b						
	d) None of the above						
6.	Consider the following reactions,						
	A: $H_2CO_3(aq) + HSO_4(aq) \leftrightarrow H_2SO_4(aq) + HCO_3(aq)$						
	B: HF (aq) + Cl <sup>-</sup> (aq) $\longrightarrow$ HCl(aq) + F <sup>-</sup> (aq)						
	C: $HF(aq) + NH_3(aq) \longrightarrow NH_4^+ + F^-(aq)$						
	D: $HSO_4^-(aq) + CN^-(aq) \longrightarrow HCN(aq) + SO_4^{2-}(aq)$						
	Reactions proceeding to the right are						
	a) A,B b) C,D c) A,C d) B,D						
7.	Conjugate base of $H_2$ is						
_	a) $H^+$ b) $H_3^+$ c) $H^-$ d) $H_3^-$						
8.	Conjugate base of $HO_2$ is						
	a) $O_2^-$ (superoxide ion) b) $H_2O_2$						
	c) $O_2^{2-}$ (peroxide ion) d) $O_2^+$						

9. In the following  $[Al(H_2O)_6]^3 + HCO_3 \quad \longleftarrow \quad [Al(H_2O)_5OH]^2 + H_2CO_3$ (A) **(B)** (C) (D) Species behaving as Bronsted- Lowry acids are b) (B), (C) d) (A), (C) a) (A), (D) c) (B), (D)10. Which is Bronsted-Lowry acid as well as Arrhenius acid? **b) HCO**<sub>3</sub><sup>-</sup> c) NH<sub>3</sub> a) H<sub>2</sub> d)  $NH_2^-$ 11. Following reaction,  $NH_4^+(aq) + CN^-(aq)$  $HCN(aq) + NH_3(aq)$ Proceeds in a) Forward direction b) backward direction c) In both sides equally d) can't be predicted

12.Out of the following, amphiprotic species are

I: $H_2PO_2^-$	II: $HPO_3^{2-}$
III: HCO <sub>3</sub> -	IV: $CH_3 CO_2^-$
V: HPO <sub>4</sub> <sup>2-</sup>	
a) I,II,III,IV	b) I,V
c) III,V	d) II,III,IV

13. What are th correct statement(s) about H<sup>-</sup> (hydride)

- a) It is Bronsted-Lowry base of H<sub>2</sub>
- b) It resembles halide
- c) It is isoelectronic of He
- d) All the above are correct statements

### 14.Consider the following reactions

I:  $[Al(OH)_3 (H_2 O)_3] + OH^- \qquad [Al(OH)_4 (H_2 O)_2^-] + H_2O$ II.  $[Al(OH)_3 (H_2 O)_3] + H_3O^+ \qquad [Al(OH)_2 (H_2 O)_4^-] + H_2O$ Select the correct statement. a) **X** is an acid in L and base in **H** 

- a) X is an acid in I and base in II
- b) X is a base in I and acid in II
- c) X is a base in I and II both
- d) X is an aid in I and II both

15.If the following proceed in forward side

HNO<sub>2</sub> +HF  $\longleftrightarrow$  H<sub>2</sub>F<sup>+</sup> + NO<sub>2</sub><sup>-</sup> CH<sub>3</sub>COOH + HF  $\iff$  F<sup>-</sup> +CH<sub>3</sub>COOH<sub>2</sub><sup>+</sup> `H<sub>2</sub>O +CH<sub>3</sub>COOH  $\iff$  H<sub>3</sub>O<sup>+</sup>CH<sub>3</sub>COO<sup>-</sup>

Then increasing order of acid strength is

### a) $H_2O < CH_3COOH < HF < HNO_2$

- b)  $HNO_2 < HF < CH_3COOH < H_2O$
- c)  $HNO_2 < HF < H_2O < CH_3COOH$
- d)  $HNO_2 < CH_3COOH < HF < H_2O$

16.Select the correct statement(s) about H<sub>2</sub>PO<sub>4</sub><sup>-</sup> and NH<sub>4</sub><sup>+</sup>

- a) Both behave as Bronsted- lowry acid
- b) H<sub>2</sub>PO<sub>4</sub><sup>-</sup> is a lewis –baseas well as Bronsted- lowry base
- c)  $H_2PO_4^-$  is amphiprotic

### d) All the above are correct statemanets

- 17.In the interaction of B(CH<sub>3</sub>)<sub>3</sub> with N(CH<sub>3</sub>)<sub>3</sub>
  - a) B(CH<sub>3</sub>)<sub>3</sub> is a lewis base and N(CH<sub>3</sub>)<sub>3</sub> is a lewis acid
  - b) B(CH<sub>3</sub>)<sub>3</sub> is a lewis acid and N(CH<sub>3</sub>)<sub>3</sub> is a lewis base
  - c) Both are lewis acids
  - d) Both are lewis bases
- 18.A carbocation (as  $CH_3$ ) is
  - a) A lewis acid b) an electrophilic reagent

c) Both a&b

d) None of these

- 19. Which is/are Lewis acids out of CH<sub>3</sub> (carbon ion ), CH<sub>5</sub>(carbonium ion ), CH<sub>3</sub> (carbonium ion)?
  - a) CH<sub>5</sub>, CH<sub>3</sub> (carbenium ion)? b) CH<sub>3</sub>, CH<sub>3</sub> c) CH<sub>3</sub> d) CH<sub>5</sub>
- 20. Which behaves as a lewis base as well as Bronsted-Lowry base?
  - a) Carbonium ion b) carbenium ion

c) carbanion d) All of these

- 21. Which s strongest acid?
  - a) **HClO**<sub>4</sub> b)  $H_2SO_4$  c)  $HNO_3$  d) HCl

22. Relative strength cannot be compared in aqueous solution of the following pair a) HClO<sub>4</sub> and HBr b)HClO<sub>4</sub> and HNO<sub>3</sub> c)  $H_2 SO_4$  and  $HNO_3$  d) All of these 23.CH<sub>3</sub>COOH (a weak acid )behaves as strong acid in a) HF b)  $NH_3$ c)  $H_2O$ HNO<sub>3</sub> 24.Select the correct acid –base equilibrium a)  $HNO_3 + HF \iff$  $H_2NO_3^+ + F^$ b)  $HNO_3 + HF \iff$  $H_2F^+ + NO_3^ CH_3COOH_2^+ + NH_2^$ c)  $CH_3COOH + NH_3$  $\longleftrightarrow$ d) HF +H<sub>2</sub>O  $\iff$  $H_2F^+ + OH^-$ 25. Consider following statements I: CH<sub>3</sub>COOH(a weak acid) behaves as a strong acid in NH<sub>3</sub> II: HNO<sub>3</sub> (a astrong acid) behaves as a base in HF III: H<sub>2</sub>SO<sub>4</sub> dissociates to a very small extent in glacial CH<sub>3</sub>COOH. Select correct alternate for these statements d) I,II, III a) I, III b) II, III c) I,II 26.100 ml of I M HCl is mixed with 50ml of 2 M HCl. Hence,  $[H_3O^+]$  is a) 1.00M b)1.50M c)1.33M d)3.00M 27.[Cl<sup>-</sup>] in a mixture of 200ml of 0.01M HCl and 100 ml of 0.01 M BaCl<sub>2 is</sub> a) 0.01 M b) 0.0133 M c) 0.03M d) 0.02M 28.B;ue litmjus turns red in the following mixture of acid and base a) 100ml of  $1 \times 10^{-2}$  M H<sub>2</sub>SO<sub>4</sub> + 100Ml of  $1 \times 10^{-2}$  M Ca(OH)<sub>2</sub> b) 100ml of  $1 \times 10^{-2}$  M HCl + 100Ml of  $1 \times 10^{-2}$  M Ba(OH)<sub>2</sub> c) 100ml of 1 ×10<sup>-2</sup> M H<sub>2</sub>SO<sub>4</sub> + 100Ml of 1 ×10<sup>-2</sup> M NaOH d) 100ml of  $1 \times 10^{-2}$  M HCl + 100Ml of  $1 \times 10^{-2}$  M NaOH 29.[H<sub>2</sub>O] in M H<sub>2</sub>SO<sub>4</sub> a) **2M** b) 1M c) 0.2M d) 0.1M 30.HCl exists as **a)** Cl **b**)  $H_2O^+$ c) HCl d) H<sub>3</sub>O<sup>+</sup>Cl 31.Select aprotic solvnts out of Water, benzene, methylamine, carbntetrachloride Ι Π III IV a) I b) II.IV c) II,III,IV d) I,III,IV 32.Strongest acid and strongest base pair is a)  $H_3O^+, H_2O$ **b**)  $H_{3}O^{+}$ ,  $OH^{-}$ c)  $H_2O, H_3O^+$ d)  $OH^{-,}H_{3}O^{+}$ 

33. Autoprotolysis constant of NH3 is

C)[NH4] [NH2]	$d)[NH_4]/[NH_2]$

34. Autoprotolysis constant of a weak acid HA is  $4 \times 10^{-8}$  hence,

[A] is

- a)  $2 \times 10^{-4} M$  b)  $0.25 \times 10^{-6} M$
- c)  $16 \times 10^{15}$  M d)  $4 \times 10^{-8}$  M
- 35.At -50°C autoprotolysis of NH<sub>3</sub> gives  $[NH_4^+]=1\times 10^{-15}$  M hence,

autoprotolusis constant of NH3 is

- a)  $\sqrt{1 \times 10^{-15}}$  b)  $\sqrt{1 \times 10^{-30}}$
- c)  $1 \times 10^{-15}$  d) $2 \times 10^{-15}$

**PH-Scale** 

36.pH of 10<sup>-8</sup> N NaOH is

**a**) 8.0 b) 6.0 c) 6.98 **d**) 7.02

37.Milliequivalent of  $Ba(OH)_2$  present in 100ml solution to have pH = 13, is

a) 10<sup>-2</sup> b) 10 c) 0.1 d)0.01

38.At a temperature under high pressure

$$K_w(H_2O) = 1 \times 10^{-10}$$

A solution of pH 5.4 under these conditons is said to be

a) Acidic **b) basic** c) neutral d) amphoteric

39.A weak monobasic acid is 0.1% ionised at 0.1M hence, its pH is

a) 2 b) 3 c)4 d)5

40. Which has maximum pH?

a) 0.01 M H<sub>2</sub>SO<sub>4</sub> b) 0.01 M HCl

c) 0.01 M Ca(OH)<sub>2</sub> d) 0.01 M NaOH

41.In which case change in pH is maximum?

a) 1 ml of pH = 2 is diluted to 100 ml

b) 0.01 mol of NaOH is added into 100ml of 0.01M NaOH solution

c) 100ml of  $H_2O$  is added into 900ml of  $10^{-6}$  M HCl

# d) 100ml of pH =2 solution is mixed with 100ml of pH =12

42.A weak monoacid base has pH = 10 at 0.01M% ionisation of base is

**a**) 0.01% **b**)0.001% **c**) 0.0001% **d**) **1.0%** 

43.100ml of solution of pH =6 is diluted to 1000ml resulting solution has pH

a) 7.0 **b) 6.7** c)7.3 d)6.4

44.In a basic solution pH is 12.3. Hence[OH<sup>-</sup>] is

a)  $2 \times 10^{-2} M$  b) $5 \times 10^{-13} M$ 

c) $3 \times 10^{-12}$  M d)) $4 \times 10^{-16}$  M

45. When you calculate pHof a solution  $[H_3O^+]$  from  $H_2O$  is required in the

following

a) 10<sup>-5</sup> M HCl b)10<sup>-5</sup> M NaOH

c)10<sup>-8</sup> M NaOH d)10<sup>-6</sup> M Ca(OH)<sub>2</sub>

### **Ostwald's Dilute Law**

46.The K<sub>a</sub> value for the acid HA is  $1.0 \times 10^{-6}$ . What is the vaue of K for the

reaction?

A-  $+H_3O \iff HA + H_2O$ b) 1.0×10<sup>8</sup> c) 1.0×10<sup>-3</sup> **a)**  $1.0 \times 10^{-8}$ d) 1.0×10<sup>6</sup>  $47.pK_b$  of aq. NH<sub>3</sub> is 1.74, hence pH of 0.01 M NH<sub>3</sub> solution is d)12.00 b)10.63 c) 2.00 a) 3.37 48.CH<sub>3</sub>COOH is 2.0% ionised ( $K_a = 1.8 \times 10^{-5}$ ), hence its molar concentration is b)0.02 M c) 3.6×10<sup>-5</sup> M a) 0.045 M d)0.090 M 49.A weak monobasic acid is 0.01% ionised when its concentration is 1M. Hence, [OH<sup>-</sup>] is d)  $1 \times 10^{10} \, \text{M}$ a) 1×10<sup>-4</sup> M b)1×10<sup>-10</sup> M c)  $1 \times 10^4 \, \text{M}$ 50.% ionisation of a weak acid is 1% at 1M, hence % solution is ...... At 4Ma) 4% b)5% c)0.5% d)0.2%

51.A mixture of weak acid is 0.1 M in HCCOH ( $K_a = 1.8 \times 10^{-4}$ ) and 0.1 M in HOCN ( $K_a = 3.3 \times 10^{-4}$ ) hence [ $H_3O^+$ ] is a) 7.14 ×10<sup>-3</sup> M c)0.20M d) $4.1 \times 10^{-3}$ b)  $4.1 \times 10^{-4} \,\mathrm{M}$ 52. At 25°C,  $[H_3O^+] = 1 \times 10^{-7}$  M in water, hence K<sub>a</sub> is a) 1×10<sup>-14</sup> b)5.55×10<sup>-15</sup> c) 1.8×10<sup>-16</sup> d) 55.5×10<sup>-10</sup> 53. What is the value of  $K_w$  in 0.01 M NaOH? d) 1×10<sup>-14</sup> **a)** 1×10<sup>-15</sup> b)1×10<sup>-13</sup> c)1×10<sup>-16</sup> 54.0.1M solution of CH<sub>3</sub>COOH should be diluted to ------ times so that pH is doubled. b) 5.55×10<sup>4</sup> times a) Four times c)  $5.55 \times 10^{6}$  times d)  $10^{-2}$  times 55. $[H_3O^+]$  in 0.1 M H<sub>2</sub>SO<sub>4</sub> at two stages  $H_2SO_4 \leftrightarrow H^+ + HSO_4^ HSO_4^- \leftrightarrow H^+ + SO_4^{2-}$  are a) 0.1M, 0.1 M b) 0.1M > 0.01M

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d) 0.1M<0.1M

C) >0.1M, >0.1M

# **Buffer solution**

56. The pH of blood is 7.40. what is the ratio of  $[HPO_4^{2-}]/[H_2PO_4^{-}]$  in the blood? ( $pK_a(H_2PO_4) = 7.10$ ) a) **2:1** b) 1.:2 c) 3:1 d)1:3 57.20 ml of 0.2M NaOH is added to 50ml of 0.2M CH<sub>3</sub>COOH, Hence (pHpK<sub>a</sub>) is b) $\log_3^2$  c)  $\log 2$ a)  $\log_{\frac{3}{2}}^{3}$ d) 2log 2 58.50.0 ml of 0.3 M HCl is mixed with 5.ml of 0.4M NH<sub>3</sub> solution. If pK<sub>a</sub> of  $NH_4^+$  is 9.26, pH of the mixture is a) 5.22 b)1.30 c) 8.78 d)12.70 59.pH of a solution made by mixing 50 of 0.2M NH<sub>4</sub>Cl and 75ml of 0.1 M NaOH is  $[pK_a \text{ of } NH_39(aq) = 4.74]$ b) 13.0 c) 4.26 **a)** 7.02 d)9.74 60.pH of a mixture which is 0.1M in CH<sub>3</sub>COOH and 0.05M in (CH<sub>3</sub>COOH)<sub>2</sub> Ba is  $[pK_a \text{ of } CH_3COOH = 4.74]$ b) 5.04 a) 4.74 c) 4.44 d) 7.00 61.We have acidic buffer of CH<sub>3</sub>COONa and CH<sub>3</sub>COOH. One or more of the following operations will not change pH I. Diluting the mixture ten times

- II. Adding some HCl
- III. Adding some NaOH

- IV. Adding equal moles of CH<sub>3</sub>COONa and CH<sub>3</sub>COOH into the buffer Select correct alternate.
- a) I, II,III,IV b)II,III c)**I,IV** d) II,IV

62. Which buffer solution has maximum pH?

- a) Mixture which is 0.1M in CH<sub>3</sub>COOH and 0.1 M in CH<sub>3</sub>COONa [pK<sub>a</sub>(CH<sub>3</sub>COOH)=4.74]
- b) Mixture which is 0.2M in CH<sub>3</sub>COOH and 0.2 M in CH<sub>3</sub>COONa
- c) Mixture which is 0.1M in NH<sub>4</sub>Cl and 0.1 M in NH<sub>4</sub>OH [ $pK_a$

 $(NH_4^+)=9.26]$ 

d) All the solutions have equal pH which is 4.74

63.The pK<sub>a</sub> of an amino acid is 9.15. at what pH amino acid is 5% dissociated?

a) 9.15 b) 4.85 c) 9.44 d)**7.87** 

64. In what volume ratio should you mix 1.0M solution of NH<sub>4</sub>Cl and NH<sub>3</sub>

to produce buffer solution of pH  $9.80?[pK_a(NH_3) = 4.74]$ 

a) **1:3.5** b) 3.5:1 c) 2:1 d)1:2

65.A weak acid HA has degree of dissociation X, thus  $(pH - pK_a)$  is

**a)**  $P_x$  **b)**  $P_{1-x}$  **c)**  $P_{1-x} + P_x$  **d)**  $P_{1-x} - P_x$ 66.pH of blood is

a) **7.4** b) 6.4 c) 8.0 d)7.0

 $67.pK_{a1}$  of carbonic acid in blood at body temperature (37°C) is 6:1, hence ratio  $[HCO_3^-]/[H_2CO_3]$  is approximately a) 1.3:1 b)1:1.3 c) 20:1 d)1:20 68. Acidosis may cause-----of the pH of blood a) Increase **b)** decrease c) no change d)is not related term 69.Important diagnosic analysis in the blood is a)  $[H_2PO_4^-]/[H_2PO_{42}^{2-}]$ b) [HCO<sub>3</sub><sup>-</sup>]/[CO<sub>2</sub>] d) [PO<sub>3</sub><sup>2-</sup>]/ [HCO4<sup>2-</sup>] c)  $[CO_3^{2-}]/[HCO_3^{2-}]$ 

70.Oxygen from inhaled air combines with haemoglobin, and oxygenated haemoglobin ionises releasing a proton which is removed in the following reactions

- a)  $H^+ + CO_2 \leftrightarrow HCO_3^-$
- b)  $HCO_3^- + H_2O \leftrightarrow H_3O^+ + CO_3^{-2-}$
- c)  $HCO_3^- + H^+ \leftrightarrow H_2CO_3$
- d) in all the above



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### **Common Ion Effect**



a) 0.005	b)0.0025	c) 0.01	d)0.002
75.[Ag <sup>+</sup> ] in saturate	ed AgCl in presence	e of 1M KCl[ K(Ag	Cl) = $1 \times 10^{-10}$ ] is
a) 1×10 <sup>-5</sup> M	b) 1×10 <sup>-20</sup> M	c)1×10 <sup>-10</sup> M	d) 2×10 <sup>-10</sup> M

76.Degree of ionisation of 1 M HCOOH is decreased to a maximum extent

in presenceof	
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- a) I M HCHO b) 1 M NaOH
- c) 1 M HCOONa d) equally for all

77.Solubility of  $Ag_2CrO_4$  is decreased in presence of

a) AgNO<sub>3</sub> b)AgCl c)BaCrO<sub>4</sub> d)PbCrO<sub>4</sub>

78.Blood pH is controlled by concentration of  $H_2CO_3$  and  $HCO_3^-$ . in

presence of NaHCO3, pH of blood is

- a) **Increased** b) decreased
- c) no change d) statement is wrong

79.A weak acid (HA) is 4% ionised at 1 M percent ionisation is 1%

presence of A<sup>-</sup> of

a) 0.04M b)0.16M c)0.02M d)0.10M

80.[OH<sup>-</sup>] in a solution prepared by mixing equal volumes of 0.M methyl

amine (CH<sub>3</sub>NH<sub>2</sub>,  $K_b = 3.7 \times 10^{-4}$ ) and 0.60 M CH<sub>3</sub>NH<sub>3</sub><sup>+</sup>Cl<sup>-</sup>

**a)**  $3.7 \times 10^{-4}$  M b)7.4  $\times 10^{-4}$  M c)  $3.7 \times 10^{-2}$  M d)1.85  $\times 10^{-4}$  M

### **Solubility Product**

- 81. Expression for  $K_{sp}$  of  $Hg_2 (NO_3)_2$  is
  - a)  $[Hg^+]^2 [NO_3^-]^2$  b) $[Hg_2^+]^2 [NO_3^-]^2$
  - c)  $[Hg^{2+}][NO_3^-]^2$  d)  $[Hg_2^{2+}]^2 [NO_3^-]^2$

82. Solubility of BaSO<sub>4</sub> in aqueous solution is  $1 \times 10^{-5}$  M. hence, solubility in

- $0.1M \; BaCl_2 is$
- a)  $1 \times 10^{-1}$  M b)  $1 \times 10^{-9}$  M c)  $1 \times 10^{-4}$  M d)  $1 \times 10^{-5}$  M

83.Molar solubility of Al(OH)2 is increased in presence of

- a) NaOH b) HCl
- c) Both a&b d) None of these

84.A saturated solution prepared by dissolving  $Ag_2CO_3$  in water has  $[Ag^+] =$ 

- $2.56{\times}10^{\text{--}4}\,M,$  its  $K_{sp}\,is$
- a) **8.4**×**10**<sup>-12</sup> b) 1.68×10<sup>-13</sup>
- c)  $6.6 \times 10^{-8}$  d) $1.6 \times 10^{-2}$

85.K<sub>sp</sub> (BaSO<sub>4</sub>) is  $1.1 \times 10^{-10}$ . in which case is BaSO<sub>4</sub> precipitated?

# a) 100 ml of $4 \times 10^{-3}$ M BaCl<sub>2</sub> + 300ml of $6.0 \times 10^{-4}$ M Na<sub>2</sub>SO<sub>4</sub>

- b) 100 ml of  $4 \times 10^{-4}$  M BaCl<sub>2</sub> + 300ml of  $6.0 \times 10^{-8}$  M Na<sub>2</sub>SO<sub>4</sub>
- c) 300 ml of  $4 \times 10^{-4}$  M BaCl<sub>2</sub> + 300ml of  $6.0 \times 10^{-8}$  M Na<sub>2</sub>SO<sub>4</sub>
- d) In all cases

86.K<sub>sp</sub> of Al(OH)<sub>3</sub> is  $1.0 \times 10^{-15}$ . pH of the saturated solution is about

**a**) 5.0 b) 9.0 c) 4.1 **d**)**10.4** 

 $87.K_{sp}\, of\, H_2S$  is  $1{\times}10^{-22}$  . [S^2-] in a buffer of pH 6 is

a)  $1 \times 10^{-16} M$  b) $1 \times 10^{-12} M$  c) $1 \times 10^{-10} M$  d) $1 \times 10^{-8} M$ 

88.Ksp of CdS is  $8.0 \times 10^{-27}$  and that of H<sub>2</sub>S is  $1 \times 10^{-22}$ .  $1 \times 10^{-14}$  M, CdCl<sub>2</sub>

solution is precipitated on passing H<sub>2</sub>S when pH ia about

a) 4 b) 6 c) **5** d)7

89.K<sub>sp</sub> of Mg(OH)<sub>2</sub> is  $1.8 \times 10^{-11}$  at 30°C. Its molar solubility is.....at pH =12

a)  $1.8 \times 10^{-11} \text{ M}$  b) $1.8 \times 10^{-9} \text{ M}$  c) $1.8 \times 10^{-54} \text{ M}$  d) $1.8 \times 10^{-7} \text{ M}$ 

90.In group III analysis buffer used to precipitate cations as hydroxide is

- a) NH<sub>4</sub>Cl +NH<sub>4</sub>OH
- b)  $HCO_3 + CO_2$
- c)  $CH_3COOH + C_6H_5COONa$
- d)  $C_6H_5COOH + C_6H_5COONa$

### Hydrolysis

91.100 ml of 0.02 M benzoic acid ( $pK_n = 4.2$ ) is titrated using 0.02M NaOH.

pH after 50ml and 100ml of NAOH have been added are

a) 3.50, 7 b)4.2, 7 c)4.2, 8.1 d)4.2, 8.25

92.pH of 0.01 M aqueous solution of NaX, NaY and NaZ are 8,9 and 10 respectively. Strongest acidof HX, HY and HZ is

a) **HX** b)HY c) HZ d) Cant be predicted

93.40 ml of 0.025M solution of the protonated form of the amino acid phenyl alanine ( $H_2A^+$ ) is treated with ml of 0.1M NaOH. pH at this stage is

 $(pK_{a1}=1.82, pK_{a2}=9.13 \text{ of } H_2A^+)$ 

a) **5.48** b)7.00 c)1.82 d)9.13

94.pH of 0.05M calcium acetate solution ( $pK_a=4.74$ ) is

a) 8.72 b) 8.87 c)7.00 d)1.30

95. Which of the aqueous solution turns blue litmus red?

**a)** NH<sub>4</sub>Cl **b)** Al(H<sub>2</sub>O)<sub>6</sub><sup>3+</sup> **c)** Fe(H<sub>2</sub>O)<sub>6</sub><sup>3+</sup> **d)** all of these

96.pK<sub>a1</sub>, pK<sub>a2</sub>, and pK<sub>a3</sub> of  $H_3PO_4$  are respectively x,yand z. pH of 0.01 M

Na<sub>2</sub>HPO<sub>4</sub> solution is

a) 2 b)  $\frac{x+y}{2}$  c)  $\frac{y+z}{2}$  d)  $\frac{x+y+z}{2}$ 

97.100ml of 0.01 M CH<sub>3</sub>COOH is titrated with 0.01 MKOH. At wat point

pH is maximum?

- a) After addition of 100ml of KOH
- b) After addition of 75ml of KOH
- c) After addition of 50ml of KOH
- d) After addition of 25ml of KOH

 $98.pK_b$  of  $NH_3$  is 4.74 and  $pK_b$  of  $A^-$ ,  $B^-$ , and  $C^-$  are 4,5, and 6 respectively.

Aqueous solution of 0.01 M ha pH in the increasing order

a)  $NH_4A < NH_4B < NH_4C$ 

# b) $NH_4C < NH_4B < NH_4A$

- c)  $NH_4C < NH_4A < NH_4B = 7$
- d) All have equal pH being salt of weak acid and weak base

 $99.pK_1$  and  $pK_2$  of  $H_2$  CO<sub>3</sub> are respectively 6.38 and 10.26. pH of 1M and

0.1M NaHCO3 are respectively

a) 8.32, 7.32 b) 7.32, 8.32 c) 8.32, 8.32 d) 7.32, 7.32

100. Degree of hydrolyss of the following is independent od

concentration

- I. NH<sub>4</sub>CN
- II. NH<sub>4</sub>HCO<sub>3</sub>
- III. NaHS
- IV. CH<sub>3</sub>NH<sub>3</sub>Cl

a) I,II,III,IV b)I,IV c) I,III,IV d) I,II,III

# Acid – base Titration and indicator

- 101. Following are some of the certain facts of Ostwalds theory of acidbase indicators
  - A. Ionised and unionised foms have different colours
  - B. Colour change is indicated at the end point when unionised form changes to ionised form due to change in pH

C. Benzenoid form changes to quinonoid form and vice-versa due to change in pH

Select the correct facts.

- a) A,B,C b) A,C c) A,B d) B, C
- 102. Select the correct statements
  - a) Methyl orange is red in alkali solution and yellow in acid solution
  - b) Phenolphthalein is pink in alkali solution and colourless in acid solution
  - c) Both a&b
  - d) None of the above
- 103. Indicator can exist in two forms depending on the medium



Select the correct statements.

- a) Methyl orange assumes form (A) in alkali solution and form (B) in acid solution
- b) Phenolphthalein assumes form (A) in acid solution and form (B) in alkaline solution
- c) Thymoohthalein assumes form (A) ain acid solution and form (B) in alkali solution
- d) All Of the above

104. For any weak acid indicator HIn ionising as  $HIn \leftrightarrow H^+ + In^-$ Colour the unionised form (HIn) is observed when a)  $\frac{In}{HIn} = \frac{1}{10}$  b) pH =pK<sub>n</sub>-1 c) Both a&b d) None of these

105. In the above case colour of the ionised form (In<sup>-</sup>) is observed when

a)  $\frac{In}{HIn} = 10$  b) pH =pK<sub>a</sub>+1 c) both a&b d) None of thes

106. Select the correct statements about indicators.

- a) Near the equivalence point  $pH = pK_a$  for weak acid indicator and  $pH = (14 pK_b)$  for weak base indicator
- b) Most indicators have a transition range of two pH points
- c) Methyl orange assumes quinonoid form in acid solution
- d)All of the above statements are correct
- 107.  $pK_a$  of the indicator is. At the equivalence point pH isa) 5b)10c)7.5d)7.0
- 108. An indicator is a weak acid and pH range of its colour s 3 to5. If the neutral points of the indicator lies in the centre of the hydrogen ion concentration corresponding to given pH range then pH at the equivalence points is

a) **3.3** b)4.0 c)7.0 d)5.0

109. In the titration of weak acid with strong base which indicator cant be used

a) Methyl orange b) phenolphthalein

c)both a&b d) none of these

110. In the titration of weak base with strong acid indicator cant be useda) Methyl orangeb) phenolphthalein

c)both a&b d) none of these

111. Suitable indicator to be used in the titration of weak acid versus strong base is

a)**phenolphthalein** b)thymophthalein c)botha&b d)None of these

112. Titration curve if a strong base is titratd with stron acid is



113. Phenolphthalein cant be used in the titration ofa) HCl with NaOHb) CH<sub>3</sub>COOH with NaOH

c) HCl with NH<sub>4</sub>OH d)CH<sub>3</sub>COOH with NH<sub>4</sub>OH

- 114. In the titration of Na<sub>2</sub>CO<sub>3</sub> with HCl, indicator used isa) Phenolphthalein **b)methyl red** 
  - c) both a&b d) none of these
- 115. 100ml of 0.1M HCl is titrate using 0.1M NaOH using phenolphthalein indicator. pH after 50ml of NaOH has been added is
  a) 7.0 b)-1.48 c)6.0 d)1.48