# びdoubtnut 

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## CHEMISTRY

## BOOKS - CENGAGE CHEMISTRY (ENGLISH)

## IONIC EQUILIBRIUM

## Solved Examples

1. Write the conjugate bases for the following Bronsted acids
(a) HF (b) $\mathrm{H}_{2} \mathrm{SO}_{4}$ (c) $\mathrm{HCO}_{3}^{\Theta}$

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2. Wirte the conjugate bases for the following :
$\Theta$
a. $\mathrm{NH}_{2}$ b. $\mathrm{NH}_{3}$ c. $\mathrm{HCO}_{3}^{\Theta}$
3. The species $\mathrm{H}_{2} \mathrm{O}, \mathrm{HCO}_{3}^{\Theta}, \mathrm{HSO}_{4}^{\Theta}, \mathrm{NH}_{3}$ can act both as Bronsted acids and bases. For each case write the corresponding conjugate acid.

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4. Classify the following species into Lewis acids and Lewis bases and show how these act as such:
(a) $\mathrm{HO}^{-}$, (b) $\mathrm{F}^{-}$, (c ) $\mathrm{H}^{+}$, (d) $\mathrm{BCl}_{3}$

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5. In the reaction of $\mathrm{BeF}_{2}$ with $2 F^{\Theta}$ to form $\mathrm{BeF}_{4}^{-2}$, which reactant is the Lewis acid and which is the Lewis base?

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6. Write the conjugate bases of the following acids:
$\oplus$
a. $\mathrm{N}_{2} \mathrm{H}_{5}$ b. $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$

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7. Write the conjugate base of the following acid:
$\oplus$
$\mathrm{N}_{2} \mathrm{H}_{5}$

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8. Liquid $\mathrm{NH}_{3}$, like water, is an amphiprotic solvent. Write the equaiton for the auto-ionisation of $\mathrm{NH}_{3}$.

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9. Aniline $\left(\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{NH}_{2}\right)$ is a weak organic base in aqueous solution.

Suggest a solvent in which aniline would become a strong base.
10. The concentration of hydrogen ion in a sample of soft drink is $2.8 \times 10^{-4} \mathrm{M}$. What is its $p H$ ?

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11. Calculate the pH of the following solutions:
a $10^{-2} \mathrm{MHCl}$
b $10^{-3} \mathrm{MH}_{2} \mathrm{SO}_{4}$
c $0.2 \times 10^{-2} \mathrm{MNaOH}$
d $0.3 \times 10^{-3} \mathrm{MCa}(\mathrm{OH})_{2}$

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12. Calculate the concentration of hydrogen ion in the acidic solution with pH
a. 4.3
13. Calculate the concentration of OH in the solution of base with pH
a. 10.4771

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14. Calculate the $p H$ of the following mixtires of strong acids, strong bases, and combination of both:
a. 500 mL of $0.1 \mathrm{MHCI}+200 \mathrm{~mL}$ of $0.1 \mathrm{MH}_{2} \mathrm{SO}_{4}+300 \mathrm{~mL}$ of $0.2 \mathrm{MHNO}_{3}$
b. 100 mL of $0.1 \mathrm{MHCI}+100 \mathrm{~mL}$ of $0.2 \mathrm{MH}_{2} \mathrm{SO}_{4}+100 \mathrm{~mL}$ of $0.1 \mathrm{MHNO}_{3}$ and 700 mL of $\mathrm{H}_{2} \mathrm{O}$
c. 500 mL of $0.1 \mathrm{MNaOH}+100 \mathrm{~mL}$ of $0.1 \mathrm{MCa}(\mathrm{OH})_{2}+400 \mathrm{~mL}$ of 0.2 MKOH d. 100 mL of $0.1 \mathrm{MNaOH}+200 \mathrm{~mL}$ of $0.1 \mathrm{NCa}(\mathrm{OH})_{2}+200 \mathrm{~mL}$ of 0.1 MKOH and 500 mL of $\mathrm{H}_{2} \mathrm{O}$
e. 100 mL of $0.1 \mathrm{MHCI}+300 \mathrm{~mL}$ of $0.1 \mathrm{MH}_{2} \mathrm{SO}_{4}+100 \mathrm{~mL}$ of $0.3 \mathrm{MBa}(\mathrm{OH})_{2}$ and volume was made to $1 L$ by adding water
f 500 mL of $0.1 \mathrm{MHCI}+100 \mathrm{~mL}$ of $0.1 \mathrm{NH}_{2} \mathrm{SO}_{4}+400 \mathrm{~mL}$ of $0.1 \mathrm{MCa}(\mathrm{OH})_{2}$
g 8 g of $\mathrm{NaOH}+680 \mathrm{~mL}$ of $\mathrm{MHCI}+10 \mathrm{~mL}$ of $\mathrm{H}_{2} \mathrm{SO}_{4}$, (specific gravity 1.2, $49 \% \mathrm{H}_{2} \mathrm{SO}_{4}$ bu mass). The total volume of the solution was made to $1 L$ with water.
h. 37.0 g of $\mathrm{Ca}(\mathrm{OH})_{2}+360 \mathrm{~mL}$ of $1 \mathrm{MHCI}+10 \mathrm{~mL}$ of $\mathrm{H}_{2} \mathrm{SO}_{4}$ (density
$=1.4,49 \% \mathrm{H}_{2} \mathrm{SO}_{4}$ by mass). The total volume of the solution was made to $1 L$ with water.

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15. a. What amount of $\mathrm{H}_{2} \mathrm{SO}_{4}$ must be dissolved in 500 mL of solution to have a pH of 2.15?

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16. Calculate the pH of solution made by mixing equal volume of three solutions having $p H=2,4$, and 6 .

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17. While calculating the pH of $10^{-7} \mathrm{MHCl}$, the common ion effect of HCl on water is considered. Why the common ion effect of HCl on water is considered?

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18. What is the pH of the following solutions:
a. $10^{-8} \mathrm{MHCl}$ b. $5 \times 10^{-8} \mathrm{MHCl}$
c. $5 \times 10^{-10} \mathrm{MHCl}$ d. $10^{-3} \mathrm{MHCl}$

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19. What is the pH of the following solutions:
$10^{-2} \mathrm{MNaOH}$

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20. Calculate the percent error in the $\left[\mathrm{H}_{3} \mathrm{O}^{\oplus}\right]$ made by neglecting the ionisation of water in $10^{-6} \mathrm{MNaOH}$ solution.

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21. The value of $K_{w}$, at the physiological temperature $37^{\circ} \mathrm{C}$ is $2.56 \times 10^{-14}$ . What is the pH at the neutral point of water at this temperature where $\Theta$ there are equal numbers of $H^{\oplus}$ and OH ions?

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22. Solution of a monobasic acid has a $\mathrm{pH}=5$. If one mL of it is diluted to 1 litre, what will be the pH of the resulting solution?

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23. The ionisation constant of $H F$ is $3.2 \times 10^{-4}$.
a. Calculate the degree of dissociation of HF present in its 1 M solution.
b. Calculate the concentration of all species present $\left(\mathrm{H}_{3} \mathrm{O}^{\oplus}, F^{\Theta}\right.$ and HF$)$ in the solution.

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24. The $p H$ of 0.1 M monobasic acid is 4.50 . Calculate the concentration of species, $H^{\oplus}, A^{\Theta}$, and $H A$ at equilibrium. Also determine the value of $K_{a}$ and $p K_{a}$ of the monobasic acid.

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25. Calculate the pH of 0.08 M solution of hypochlorous acid, HOCl . The ionization constant of the acid is $2.5 \times 10^{-5}$. Determine the percent dissociation of HOCl.
26. The pH of 0.004 M hydrazine solution is 9.7 . Calculate its ionization constant $K_{b}$ and $p K_{b}$.

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27. Determine the dergee of dissociation of $0.05 \mathrm{MNH}_{3}$ at $25^{\circ} \mathrm{C}$ in a solution of $\mathrm{pH}=11$

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28. Calculate the ionic constant of the conjugate acid of $\mathrm{NH}_{3}$. given k_(b) is $1.77 \times 10^{\wedge}-5$

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29. Prove that the dergee of dissociation of weak acid is given by:
$\alpha=\frac{1}{1+10^{p K_{a}-p H}}$
where $K_{a}$ is its dissociation constant of the weak acid.

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30. Calculate $\left[H^{\oplus}\right]$ in a soluton that is 0.1 MHCOOH and $0.1 \mathrm{MHOCN} \cdot K_{a}(\mathrm{HCOOH})=1.8 \times 10^{-4}, K_{a}(\mathrm{HoCN})=3.3 \times 10^{-4}$.

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31. Calculate $\left[H^{\oplus}\right],\left[\mathrm{C}_{3} \mathrm{H}_{5} \mathrm{O}_{3}^{\Theta}\right]$, and $\left[\mathrm{PhO}^{\Theta}\right]$ in a solution that is
$0.03 \mathrm{M}\left(\begin{array}{c}\mathrm{O} \\ \text { I। } \\ \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{C}-\mathrm{O}-\mathrm{O}-\mathrm{H}\end{array}\right)$ and $0.1 \mathrm{MPhOH} ? \mathrm{~K}_{a}$ values for $\mathrm{C}_{3} \mathrm{H}_{5} \mathrm{O}_{3} \mathrm{H}$ and
PhOH are $1.48 \times 10^{-4}$ and $1.05 \times 10^{-10}$ respectively.

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32. What is $p H$ of $7.0 \times 10^{-8} \mathrm{M}$ acetic acid. What is the concentration of un-ionsed acetic acid. $\mathrm{K}_{a}$ of $\mathrm{CH}_{3} \mathrm{COOH}=1.8 \times 10^{-5}$.

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33. The $K_{a}$ for formic acid and acetic acid are $2.1 \times 10^{-4}$ and $1.1 \times 10^{-5}$, respectively. Calculate relative strength of acids

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34. What is the pH of the solution when 100 mL of 0.1 MHCl is mixed with 100 mL of $0.1 \mathrm{MCH}_{3} \mathrm{COOH}$.

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35. Calculate $\left[H^{\oplus}\right]$ and $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$ in $10^{-3} \mathrm{M}$ solution of monobasic acid which is $4.0 \%$ ionised. What is the $p H, K_{a}$ and $p K_{a}$ of the acid.

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36. calculate $\left[H^{\oplus}\right]$ and $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$ in a $0.1 M$ solution of weak monoacitic base which is $2.0 \%$ ionised. What is the pH of solution.

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37. The pH of pure water at $25^{\circ} \mathrm{C}$ and $35^{\circ} \mathrm{C}$ are 7 and 6 , respectively. $\Theta$
Calculate the heat of formation of water from $H^{\oplus}$ and $O H$.

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38. The $p H$ of 0.05 M aqueous solution of diethy1 amine is 12.0 . Caluclate $K_{b}$.
39. What is the pH of 1 M solution of acetic acid? To what volume one litre of this solution be diluted so that pH of the resulting solution will be twice of the original value ? $\left(K_{a}=1.8 \times 10^{-5}\right)$

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40. Calculate the pH of $0.1 \mathrm{MNH}_{3}$ solution. $K_{b}=1.8 * 10^{-5}$

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41. Calculate the pH after 50.0 mL of $0.1 \mathrm{M} \mathrm{NH}_{-}(3)$ this solution is treated with 25.0 mL of 0.1 MHCI
$K_{b}$ for $\mathrm{NH}_{3}=1.77 \times 10^{-5}\left(p K_{b} \approx 4.76\right)$.

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42. What is the pH of a solution containing 0.01 MHCl ?
43. What is $\left[\mathrm{H}^{+}\right]$in $\mathrm{mol} / \mathrm{L}$ of a solution that is 0.20 M in $\mathrm{CH}_{3} \mathrm{COONa}$ and 0.10 M in $\mathrm{CH}_{3} \mathrm{COOH} ?\left(\mathrm{~K}_{a}\right.$ for $\left.\mathrm{CH}_{3} \mathrm{COOH}=1.8 \times 10^{-5}\right)$

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44. $0.1 \mathrm{MNH}_{3}$ solution is found to have a $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$ of $.133 \times 10^{-3} \mathrm{M}$.
a. What is the pH of the solution?
b. What will be the pH of the solution after 0.1 MNaOh is added to it?
c. Calculate $K_{b}$ and $p K_{b}$ for $\mathrm{NH}_{3}$ ?
d. How will NaOh added to the solution affect the extent of dissociation of $\mathrm{NH}_{3}$ ?

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45. The self ionisation constant for pure
$\mathrm{HCOOH}, \mathrm{K}=\left[\mathrm{HCOOH}_{2}{ }^{\oplus}\right]\left[\mathrm{HCOO}^{\Theta}\right]$ is $10^{-6}$ at room temperature. What percentage of HCOOH molecules are converted to $\mathrm{HCOO}^{\ominus}$ ions. The density of HCOOH iws $1.22 \mathrm{gcm}^{-3}$.

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46. Liquid $\mathrm{NH}_{3}$ dissociation to a slight extent, At a certain temp. its self dissociation constant $K_{S D C}\left(\mathrm{NH}_{3}\right)=10^{-30}$. The number of $\mathrm{NH}_{4}^{+}$ions are present per $100 \mathrm{~cm}^{3}$ of pure liquid are :

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47. Find the concentration of $\mathrm{H}^{\oplus}, \mathrm{HCO}_{3}^{\Theta}$, and $\mathrm{CO}_{3}^{-2}$ in a 0.01 M solution of carbonic acid if the pH of solution is 4.18 .
$K_{1}=4.45 \times 10^{-7}, K_{2}=4.69 \times 10^{-11}$
48. $K_{1}$ and $K_{2}$ for dissociation of $H_{2} A$ are $4 \times 10^{-3}$ and $1 \times 10^{-5}$. Calculate concentration of $A^{2-}$ ion in $0.1 \mathrm{MH}_{2} A$ solution. Also report $\left[\mathrm{H}^{+}\right]$and pH .

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49. Calculate the concentration of all species of significant concentrations
presents in $\quad 0.1 \mathrm{MH}_{3} \mathrm{PO}_{4}$ solution. If
$K_{1}=7.5 \times 10^{-3}, K_{2}=6.2 \times 10^{-8}, K_{3}=3.6 \times 10^{-13}$

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50. A solution contains $0.10 \mathrm{MH}_{2} \mathrm{~S}$ and 0.3 HCl Calculate the concentration of $S^{2-}$ and $\mathrm{HS}^{-}$ions in the solution . For $H_{2} S, K_{a_{1}}=1 \times 10^{-7}$ and $K_{a_{2}}=1.3 \times 10^{-13}$
51. Which of the following combinations of solute would result in the formation of a buffer solution.
a. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{NaOH}$ in
i. 1:1 mol ratio
ii. 2:1 mol ratio
iii. 1:2 mol ratio
b. $\mathrm{NH}_{4} \mathrm{CI}=\mathrm{NH}_{3}$ in
i. 1:1 mol ratio
ii. 2:1 mol ratio
iii. 1:2mol ratio
c. $\mathrm{HCI}+\mathrm{NaCI}$
d. $\mathrm{HCI}+\mathrm{CH}_{3} \mathrm{COOH}$
e. $\mathrm{NaH}+\mathrm{HCI}$

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52. Calculate the $p H$ of a solution made by mixing $0.1 M N H_{3}$ and
$0.1 \mathrm{M}\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4} \cdot\left(p K_{b}\right.$ of $\left.\mathrm{NH}_{3}=4.76\right)$
53. How much volume of 0.1 MHac should be added to 50 mL of 0.2 MNaAc solution to have a pH 4.91 ?

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54. i At what pH will the mixture of HCOOH and HCOONa given buffer solution of higher capacity?

$$
\left(K_{a} o f H C O O H=1.8 \times 10^{-4}\right)
$$

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55. How much ml of $0.3 \mathrm{MNH}_{4} \mathrm{OH}$ should be mixed with 30 mL of 0.2 M solution of $\mathrm{NH}_{4} \mathrm{Cl}$ to given buffer solution of $\mathrm{pH8} 8.65$ ? $(\mathrm{pKb}=4.74)$

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56. Calculate the $p H$ of the following mixtures given $K_{a}=1.8 \times 10^{-5}$ and $K_{b}=1.8 \times 10^{-5}\left(p K_{a}=p K_{b}=4.7447\right)$.
a. $50 \mathrm{~mL} 0.05 \mathrm{MNaOH}+50 \mathrm{~mL}$ of $0.1 \mathrm{MCH}_{3} \mathrm{COOH}$
b. $50 \mathrm{~mL} 0.1 \mathrm{MNH}_{4} \mathrm{OH}+50 \mathrm{~mL}$ of 0.05 MHCI

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57. What volume of strong monobasic acid of normality 10 is needed to prepare 1 L of a butter solution of $\mathrm{pH9}$, using 1 mol of $\mathrm{NH}_{3}$ and as much of strong acid needed. $\left(K_{b}\right.$ for $\left.\mathrm{NH}_{3}=1.8 \times 10^{-5}\right)\left(p K_{b}=4.7477\right)$

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58. 40 mL sample of 0.1 M solution of nitric acid is added to 20 mL of 0.3 M aqueous ammonia. What is the pH of the resulting solution? $\left(p K_{b}=4.7447\right)$
59. The base imidazole has a $K_{b}$ of $8.1 \times 10^{-8}$.
a. In what amounts should 0.02 MHCI and 0.02 M imidazole be mixted to make 100 mL of a buffer at pH 7 ?
b. If the resulting solution is diluted to $1 L$, what is the $p H$ of the diluted solution?

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60. In the titration of solution of a weak acid HA and NaOH , the pH is 5.0 after 10 mL of NaOH solution has been added and 5.60 after 20 mL NaOH has been added.

What is the value of $p K_{a}$ for HA?

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61. A definite volume of an aqueous $N / 20$ acetic acid $\left(p K_{a}=4.74\right)$ is titrated with a strongs base. It is found that 75 equal-sized drops of

NaOH added from a burette effect the complete neutralisation. Find the $p H$ when an acid solution is neutralised to the extent of $20 \%, 40 \%$, and $90 \%$, respectively.

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62. How many moles of NaOH can be added to 0.1 L of solution of $0.1 \mathrm{MNH}_{3}$ and $0.1 \mathrm{MNH}_{4} \mathrm{CI}$ without changing pOH by more than pne unit $\left(p K_{a}{ }^{\text {ofNH }}{ }_{3}=4.75\right)$ ?

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63. How many moles of HCI can be added to 1.0 L of solution of $0.1 \mathrm{MNH}_{3}$ and $0.1 \mathrm{MNH}_{4} \mathrm{CI}$ without changing pOH by more than one unit? $\left(p K_{b} o f \mathrm{NH}_{3}=4.75\right)$

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64. A buffer solution of pH value 4 is to be prepared, using $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COONa}$.How much amount of sodium acetate is to be added to 1.0 L of $M / 10$ acetic acid? $\left(K_{a}=2.0 \times 10^{-5}\right)$

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65. What will be the $p H$ if 0.01 mol of HCl is dissolved in the above buffer solution? Find the change in pH value.

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66. How will the pH be affected if 1.5 L of $\mathrm{H}_{2} \mathrm{O}$ is added to the above buffer?

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67. Calculate the pH of a buffer by mixing 0.15 mole of $\mathrm{NH}_{4} \mathrm{OH}$ and 0.25 mole of $\mathrm{NH}_{4} \mathrm{Cl}$ in a 1000 mL solution $\mathrm{K}_{b}$ for $\mathrm{NH}_{4} \mathrm{OH}=2.0 \times 10^{-5}$

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68. To 0.1 L of a decimolar solution of acetic acid, how much dry sodium acetate be added (in moles) so as to decrease the concentration of $H^{\oplus}$ ion to $1 / 10$ th of its previous value? $K_{a}=2.0 \times 10^{-5}$.

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69. The equivalent point in a titration of 40.0 mL of a sodium of a weak monoprotic acid occurs when 35.0 mL of a 0.10 MNaOH solutio has been added. The pH of the solution is 5.5 after the addition of 20.0 mL of NaOH solution. What is the dissociation constant of the acid ?

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70. Consider a buffer solution containing 0.1 mol each of acetic and sodium acetate in 1.0 L of solution, 0.01 mol of NaOH is gradually added to this buffer solution. Calculate the average buffer capacity of the solution and as well as initial and final buffer capacity. $\left[K_{a}=2 \times 10^{-5}\right] p K_{a}=4.7$

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71. The ph of blood stream is maintained by a proper balance of $\mathrm{H}_{2} \mathrm{CO}_{3}$ and $\mathrm{NaHCO}_{3}$ concentrations. What volume of $5 \mathrm{MNaHCO}_{3}$ solution, should be mixed with 10 mL sample of blood which is 2 M in $\mathrm{H}_{2} \mathrm{CO}_{3}$ in order to maintain a pH of 7.4. $\mathrm{K}_{a}$ for $\mathrm{H}_{2} \mathrm{CO}_{3}$ in blood is $7.8 \times 10^{-7}$ ?

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72. Calculate the degree of hydrolysis of a mixture of aniline and acetic acid each of them being $0.01 M . K_{a}$ of acetic acid $=1.8 \times 10^{-5}$ and $K_{b}($ aniline $)=4.5 \times 10^{-10}$. Also calculate $p H$ of the mixture.
73. 2.5mLof $2 / 5 M$ weak mono-acidic base $\left(K_{b}=1 \times 10^{-12}\right.$ at $\left.25^{\circ} \mathrm{C}\right)$ is titrated with $2 / 15 M H C I$ in water at $25^{\circ} \mathrm{C}$. Find the concentration of $H^{\oplus}$ ions at equivalence point. $\left(K_{w}=1 \times 10^{-14} a t 25^{\circ} \mathrm{C}\right)$
a. $3.7 \times 10^{-13} \mathrm{M}$ b. $3.2 \times 10^{-7} \mathrm{M}$
c. $3.2 \times 10^{-2} \mathrm{M}$ d. $2.7 \times 10^{-2} \mathrm{M}$

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74. The $p K_{a}$ of $\mathrm{CH}_{3} \mathrm{COOH}$ and $p \mathrm{~K}_{a}$ of $n \mathrm{H}_{4} \mathrm{OH}$ is 4.76 and 4.75 , respectively.

Calculate the hydrolysis constant of ammonium acetate $\left(\mathrm{CH}_{3} \mathrm{COONH}_{4}\right)$ at 298 K and also the drgree of hydrolysis and pH of its (a) 0.01 M and (b) $0.04 M$ solutions.

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75. Calculate the dergee of hydrolysis and pH of 0.02 M ammonium cyanide $\left(\mathrm{NH}_{4} \mathrm{CN}\right)$ at 298 K . $\left(K_{a}\right.$ of $H C N=4.99 \times 10^{-9}, K_{b}$ for
$\left.\mathrm{NH}_{4} \mathrm{OH}=1.77 \times 10^{-5}\right)$

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76. Calculate the $p H$ of the solutions when following conditions are provided:
a. 20 mL of $\mathrm{M} / 10 \mathrm{CH}_{3} \mathrm{COOH}$ solution is titrated with $\mathrm{M} / 10$ solution of NaOH .
i. No titration is carried out.
$\left(p K_{a}\right.$ of $\left.\mathrm{CH}_{3} \mathrm{COOH}=4.74\right)$

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77. Calculate the $p H$ of the following mixtures given $\left(p K_{a}=p K_{b}=4.7447\right)$ :
a. $50 \mathrm{~mL} 0.1 \mathrm{MNaOH}+50 \mathrm{~mL} 0.1 \mathrm{MCH}_{3} \mathrm{COOH}$
b. $50 \mathrm{~mL} 0.1 \mathrm{mNaOH}+50 \mathrm{~mL} 0.05 \mathrm{MCH}_{3} \mathrm{COOH}$
c. $50 \mathrm{~mL} 0.05 \mathrm{MNaOH}+50 \mathrm{mLO} .1 \mathrm{MCH}_{3} \mathrm{COOH}$
d. $50 \mathrm{mLO} 0.1 \mathrm{MNH}_{4} \mathrm{OH}+50 \mathrm{~mL} 0.05 \mathrm{MHCI}$
e. $50 \mathrm{mLL} 0.05 \mathrm{MNH}_{4} \mathrm{OH}+50 \mathrm{~mL} 0.1 \mathrm{MHCI}$
f. $50 \mathrm{mLO} 0.05 \mathrm{MNH}_{4} \mathrm{OH}+50 \mathrm{mLO} 05 \mathrm{MCH}_{3} \mathrm{COOH}$

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78. Which of the following ions or compounds in a solutions tends to produe an acidic, a basic, or a neutral solution.
a. i. $\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{\Theta}$ ii. $\mathrm{Na}^{\oplus}$ iii. $\mathrm{So}_{3}^{2-}$ iv. $\mathrm{F}^{\Theta}$ v. $\mathrm{NH}_{4}^{\oplus}$
b. i. $\mathrm{CH}_{3} \mathrm{COONa}$ ii. $\mathrm{ZnCI}_{2}$ iii. $\mathrm{KNO}_{3}$ iv. $\mathrm{NH}_{4} \mathrm{CI}$
c. i. NaCN ii. $\mathrm{K}_{2} \mathrm{CO}_{3}$ iii. $\mathrm{H}_{3} \mathrm{PO}_{4}$ iv. NaF
d. i. $\mathrm{NH}_{4} \mathrm{NO}_{3}$ ii. $\mathrm{Ba}_{2} \mathrm{CO}_{3}$ iii. $\mathrm{NaHSO}_{4}$ iv. NaOCI v. HOCI vi. $\mathrm{AI}\left(\mathrm{NO}_{3}\right)_{3}$

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79. Arrange the following bases in order of decreasing basicity:
$\mathrm{S}^{2-}, \mathrm{CH}_{3} \mathrm{COO}^{\Theta}, \mathrm{CN}^{\Theta}, \mathrm{NH}_{3}, F^{\Theta}$
80. Classify each of the folowing as a strong acid, string base, weak acid, and weak base:
i. NaOH ii. HF iii. $\mathrm{NH}_{4}^{\oplus}$ iv. $N H_{3}$ v. $F^{\Theta}$ vi. HI

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81. Arrange the following 0.1 M solutions in order of increasing pH :
$\mathrm{H}_{2} \mathrm{CO}_{3}, \mathrm{HBr}, \mathrm{HI}, \mathrm{NH}_{3}, \mathrm{KCN}, \mathrm{NaOH}, \mathrm{NH}_{4} \mathrm{Br}$

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82. Why the following compounds will produce acidic solution in water
i. $\mathrm{H}_{3} \mathrm{PO}_{4}$
ii. $\mathrm{CO}_{2}$
iii. $\mathrm{HNO}_{2}$
iv. $\mathrm{AlCl}_{3}$
83. Write equations \& explain why the following species act as weak bases in water solution.
i. $\mathrm{CH}_{3} \mathrm{NH}_{2}$ ii. $\mathrm{NO}_{2}^{\Theta}$ iii. $\mathrm{HPO}_{4}^{2-}$ iv. $\mathrm{CHO}_{2}^{\Theta}$

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84. Which equilibrium constant(s) or ratio of equilibrium constants should be used to calculate the pH of 1.00 L of each of the following solutions?
a. KOH b. $\mathrm{NH}_{3}$ c. $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$
d. $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+\mathrm{NaC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ e. $\mathrm{KC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$
f. $0.01 \mathrm{molHC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+0.050 \mathrm{molNaOh}$
g. $\mathrm{H}_{2} \mathrm{~S}$ h. $0.01 \mathrm{NH}_{4} \mathrm{CI}+0.50 \mathrm{molNaOH}$
i. $0.010 \mathrm{molHC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+0.10 \mathrm{molNaOH}$

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85. The salt of which one of the follwing five weak acid will be the most hydrolysed?
a. $H A: K_{a}=1 \times 10^{-8}$
b. $\mathrm{HB}: K_{a}=2 \times 10^{-6}$
c. $H C: K_{a}=3 \times 10^{-8}$
d. $H D: K_{a}=4 \times 10^{-10}$
e. $H E: K_{a}=1 \times 10^{-7}$

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86. 500 mL of 0.2 M aqueous solution of acetic acid is mixed with 500 mL of 0.2 HCI at $25^{\circ} \mathrm{C}$.
a. Calculate the degree of dissociation of acetic acid in the resulting solution and pH of the folution.
b. If 6 g of NaOH is added to the above solution determine the final pH. $\left[K_{a}\right.$ of $\mathrm{CH}_{3} \mathrm{COOH}=2 \times 10^{-5}$.

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87. 0.1 MNaOH is titrated with $0.1 \mathrm{M}, 20 \mathrm{mLHA}$ till the point. $K_{a}(H A)=6 \times 10^{-6}$ and degree of dissociation of $H A$ is neglible (small) as compared to unity. Calculate the pH of the resulting solution at the end point [Use $\log 6 \approx 0.8]$

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88. A hydrogen electrode placed in a solution containing sodium acetate and acetic acid in the ratio of $x: y$ and $y: x$, ahs electrode potential values of -1.5 and -0.5 V , respectively. What is the $p K_{a}$ value of acetic acid?

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89. The emf of the following cell is observed to be 0.118 V at $25^{\circ} \mathrm{C}$.
$\left[P t, H_{2}(1 \mathrm{~atm}) \mid H A\left(100 \mathrm{~mL} 0.1 \mathrm{M}| | H^{\oplus}(0.1 \mathrm{M})\left|H_{2}(1 \mathrm{~atm})\right| P t\right]\right.$
a. If 30 mL of 0.2 MNaOH is added to the negative terminal of battery, find the emf of the cell.
b. If 50 mL of 0.2 MNaOH is added to the negative terminal of battery, find the emf of teh cell.

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90. The freezing point of 0.20 M solution of weak acid $H A$ is 272.5 K . The molality of the solution is $0.263 \mathrm{~mol} \mathrm{Kg}^{-1}$.

Find the $p H$ of the solution on adding 0.20 M solution of NaOH . Given: $\mathrm{K}_{f}$ of water $=1.86 \mathrm{Km}^{-1}$

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91. Methy1 red has a $K_{a}=10^{-5}$. The acid form Hin is red and its conjugate base, Ind $^{\Theta}$ is yellow. Complete the following table:
$\begin{array}{llll}\mathrm{pH} & 3 & 5 & 7\end{array}$
$\left[\right.$ Ind $\left.^{\Theta}\right] /[\mathrm{HIn}] \quad-\quad-$
Colour
92. There are three acid-base indicators. Methyl orange (end point at $p H=4$ ), bromothymol blue (end point at $p H-7$ ), phenolphthalein (end point at $p H=9$ ). Which is the most suitable indicator for the following titrations?
(a). $\mathrm{H}_{2} \mathrm{SO}_{4}$ with KOH
(b). KCn with HCl
(c). $\mathrm{NH}_{3}$ with $\mathrm{HNO}_{3}$
(d). HF with NaOH

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93. A solution gives the following colours with different indicators:
a. Methyl orange $\Rightarrow$ Yellow
b. Methyl red $\Rightarrow$ Yellow
c. Bromthymol blue $\Rightarrow$ Orange

What is the pH of the solution?
94. What indicators will be suitable for the following acid-base titrations:
a. HCOOH against NaOH
b. HBr against KOH
c. $\mathrm{NH}_{4} \mathrm{OH}$ with $\mathrm{HNO}_{3}$

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95. Calculate the $p H$ at which an acid indicator with $K_{a}=1.0 \times 10^{-5}$ changes colour when the indicator is $1.00 \times 10^{-3} M$.

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96. At what $p H$ will a $1.0 \times 10^{-3} M$ solution of an indicator with $K_{b}=1.0 \times 10^{-10}$ changes colour?

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97. What indicator should be used for the titration of $1.0 \mathrm{MKBO}_{2}$ with

### 1.10MHCl?

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98. Calcualte the pH at which an indicator with $p K_{b}=4$ changes colour.

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99. Bromophenol blue is an indicator with a $K_{a}$ value of $5.84 \times 10^{-5}$. What is the percentage of this indicator in its basic form at a pH of 4.84 ?

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100. An acid-base indicator has $K_{H I n}=3.0 \times 10^{-5}$. The acid form of the indicator is red and the basic form is blue. The change in $\left[\mathrm{H}^{+}\right]$required to change the indicator from $75 \%$ red to $75 \%$ blue is :
101. Determine the solubility of (a) AgCI , (b) $\mathrm{Fe}(\mathrm{OH})_{3}$, (c) $\mathrm{Hg}_{2} \mathrm{Br}_{2}$, and (d) $\mathrm{Ag}_{2} \mathrm{SO}_{4}$ from their solubility product constants give in table. Calculate the molarities of the individual ions and also the soubities of salts in $g L^{-1}$.

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102. Calcualte the solubility of $M_{2} X_{3}$ in pure water, assuming that neither kind of ion reacts with $\mathrm{H}_{2} \mathrm{O}$. The solubility product of $M_{2} X_{3}, K_{s p}=1.1 \times 10^{-23}$.

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103. The values of $K_{s p}$ of two sparingly soluble salts $\mathrm{Ni}(\mathrm{OH})$ an d AgCN are $2.0 \times 10^{-15}$ and $6 \times 10^{-17}$ respectively. Which salt is more soluble? Explain.
104. A solution of $\mathrm{caF}_{2}$ is found to contain $10^{-4} \mathrm{MF}{ }^{\Theta}$ ions. What is the $K_{s p}$ of $c a F_{2}$ ?

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105. Let the solubilities of AgBr in water and in $0.01 \mathrm{MCaBr}_{2}, 0.01 \mathrm{MKBr}$, and $0.05 \mathrm{MAgNO}_{3}$ be $S_{1}, S_{2}, S_{3}$ and $S_{4}$, respectively. Give the decreasing order of the solubilities.

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106. The $K_{s p}$ of AgCl at $25^{\circ} \mathrm{C}$ is $1.5 \times 10^{-10}$. Find the solubility (in $g L^{-1}$ ) in an aqueous solution containing $0.01 \mathrm{MAgNO}_{3}$.

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107. The solubility of $\mathrm{BaSO}_{4}$ in water is $2.33 \mathrm{~g} 100 \mathrm{~mL}^{-1}$. Calculate the percentage loss in weight when 0.2 g of $\mathrm{BaSO}_{4}$ is washed with
a. $1 L$ of water
b. 1 L of $0.01 \mathrm{NH}_{2} \mathrm{SO}_{4} \cdot\left[\mathrm{Mw}_{\mathrm{BaSO}_{4}}=233 \mathrm{gmol}^{-1}\right]$

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108. When 15 mL of $0.05 \mathrm{MAgNO}_{3}$ is mixed with 45.0 mL of $0.03 \mathrm{MK}_{2} \mathrm{CrO}_{4}$, predict whether precipitation of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ occurs or not? $K_{s p}$ of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}=1.9 \times 10^{-12}$

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109. The concentration of $\mathrm{Ni}^{+2}$ ions in a given NiS solution is $2.0 \times 10^{-6} \mathrm{M}$.

Find the minimum $S^{2-}$ ions necessary to cause precipitation of $N i S . K_{s p}$ of $N i S=1.4 \times 10^{-14}$.
110. A solution contains 0.1 M each of $\mathrm{CaCI}_{2}$ and $\mathrm{SrCI}_{2} \cdot \mathrm{~A} 0.005 \mathrm{M}$ solution of $\mathrm{SO}_{4}^{2-}$ is slowly added to the given solution.
a. Which substance beings to precipiate first?
b. If $\mathrm{H}_{2} \mathrm{SO}_{4}$ is continuosult added, determine when will other salt be precipitated?
c. When second salt starts to precipitate, find the concentration of cation of first salt. Assume that $\mathrm{CaCI}_{2}$ and $\mathrm{SrCI}_{2}$ are $100 \%$ ionised and volume of the solution remains constant.

$$
K_{s p} o f \text { SrSO}_{4}=3.2 \times 10^{-7} \text { and } K_{s p} o f \text { CaSO }_{4}=1.3 \times 10^{-4}
$$

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111. How much the concentration of $A{ }^{\oplus}$ ions in a saturated solution of AgCI diminish if such an amount of $H C I$ is added to it that the concentration of $C I^{\Theta}$ ions in the solution becomes equal to $0.03 M$ ? Also find the amount of AgCI precipitated at the given concentration. $K_{s p}$ of $A g C I=1.8 \times 10^{-10}$.
112. Calculate the maximum possible concentration of $\mathrm{Mn}^{2+}$ in water that is saturated with $\mathrm{H}_{2} \mathrm{~S}$ (which is 0.1 M at 300 K ) and maintained at $\mathrm{pH}=3$ with HCI. The equilibrium constant (s) for dissociation of $H_{s} S$ are:
$H_{2} S \Leftrightarrow H^{\oplus}+H S^{\Theta}, K_{1}=9 \times 10^{-8}$
$H S^{\Theta} \Leftrightarrow H^{\oplus}+S^{2-}, K_{2}=1 \times 10^{-12}$ and
$K_{s p}$ of $M n S=3 \times 10^{-22}$

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113. Two weak monobasic organic acids $H A$ and $H B$ have dissociation constants as $3.0 \times 10^{-5}$ and $1.5 \times 10^{-5}$, respectively, at $25^{\circ} \mathrm{C}$. If 500 mL of $1 M$ solutions of each of these two acids are mixed to produce $1 L$ of mixed acid solution, what is the pH of the resulting solutions?

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114. Calculate the simultaneous solubility of AgSCN and $\mathrm{AgBr} . K_{s p}$ for AgSCN and AgBr are $1 \times 10^{-12}$ and $5 \times 10^{-13}$ respectively.

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115. The following equilibria are given by
rhoN $\mathrm{N}_{2}+3 \mathrm{H}_{2} \Leftrightarrow 2 \mathrm{NH}_{3}: \mathrm{K}_{1}$
$\mathrm{N}_{2}+\mathrm{O}_{2} \Leftrightarrow 2 \mathrm{NO}, \mathrm{K}_{2}$
$\mathrm{H}_{2}+\frac{1}{2} \mathrm{O}_{2} \Leftrightarrow \mathrm{H}_{2} \mathrm{O}, \mathrm{K}_{3}$
The equlibrium constant of the reaction
$2 \mathrm{NH}_{3}+\frac{5}{2} \mathrm{O}_{2} \Leftrightarrow 2 \mathrm{NO}+3 \mathrm{H}_{2} \mathrm{O}$
in terms of $K_{1}$ and $K_{3}$ is

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 solution of $p H=4,5$ and 6 are $S_{1}, S_{2}, S_{3}$, and $S_{4}$, respectively. Given the
decreasing order of their solubilities.

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117. The ionization contant of benzoic acid is $6.46 \times 10^{-5}$ and $K_{s p}$ for silver benzoate is $2.5 \times 10^{-13}$. How many times is silver benzoate more soluble in abuffer of $\mathrm{pH}=3.19$ compared to its solubility in pure water ?

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118. Write equations showing all of the equilibrium reactions occurring in aqueous solution containing following reagent:

NaCI

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119. Calculate the solubility of CoS in $0.1 \mathrm{MH}_{2} \mathrm{~S}$ and $0.15 \mathrm{MH}_{3} \mathrm{O}^{\oplus}\left(K_{\text {sp }}\right.$ of $\left.\operatorname{CoS}=3 \times 10^{-26}\right)$.

$$
\left(K_{1} \times K_{2}\left(H_{2} S\right)=10^{-21}\right)
$$

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120. Explain why $\operatorname{CoS}$ is more soluble than predicted by the $K_{s p}$.

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121. The solubility of CuS in pure water at $25^{\circ} \mathrm{C}$ is $3.3 \times 10^{-4} \mathrm{gL}^{-1}$.

Calculate $K_{s p}$ of CuS . The accurate value of $K_{s p}$ of CuS was found to be $8.5 \times 10^{-36}$ at $25^{\circ} \mathrm{C}$.

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122. Explain why CuS is more soluble than predicted by the $K_{s p}$.

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123. The solubility of $\mathrm{TI}_{2} \mathrm{~S}$ in pure $\mathrm{CO}_{2}$-free water is $6.3 \times 10^{-6} \mathrm{M}$. Assume that the dissolved $S^{2-}$ ion hydrolyses almost completely to $H S^{\Theta}$ and that the further hydrolysis to $H_{2} S$ is neglected. What is the $K_{s p} \cdot\left(K_{2}\left(H_{2} S\right)=10^{-14}\right)$.

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124. When solid $\mathrm{SrCO}_{3}$ is equilibrated with a $p H 8.60$ buffer, the solution was found to have $\left[\mathrm{Sr}^{2+}\right]=2.2 \times 10^{-4}$. What is the $K_{s p}$ of $\mathrm{SrCO}_{3}$. $\left(K_{2}\right.$ of $\left.\mathrm{H}_{2} \mathrm{CO}_{3}=4.7 \times 10^{-11}\right)$

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125. Calculate the solubility at $25^{\circ} \mathrm{C}$ of $\mathrm{CaCO}_{3}$ in a closed container containing a solution of $p$ H8.60. $\left[K_{\text {sp }}\left(\mathrm{CaCO}_{3}\right)=10^{-8}, K_{a}=4.7 \times 10^{-11}\right]$

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126. For the galvanic cell: $\operatorname{Ag}|\operatorname{AgCI}(s)| \operatorname{KCI}(0.2 M)||\operatorname{KBr}(0.001 M)| \operatorname{AgBr}(s)| \operatorname{Ag}$, calculate the EMF generated for a spontaneous process after taking into account the cell reaction at $25^{\circ} \mathrm{C}$.

$$
\left[K_{s p(\mathrm{AgCI})}=2.8 \times 10^{-10}, K_{s p(\mathrm{AgBr})}=3.3 \times 10^{-13}\right]
$$

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127. At $25^{\circ} \mathrm{C}$, after the addition of 110 mL of 0.1 NaCI solution to 100 mL of $0.1 \mathrm{NAgNO}_{3}$ solution, the reduction potentilal of a silver electrode placed in it is 0.36 V . Calculate the $K_{s p}$ of AgCI . (Given: $E^{\Theta} \mathrm{Ag} / \mathrm{Ag}^{\oplus}=-0.799 \mathrm{~V}$ ).

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$\Theta$
128. Calculate the entropy of $O H$ ion at 298 K . Given:
$\Theta$
a. $\mathrm{H}_{2} \mathrm{O} \Leftrightarrow H^{\oplus}+\mathrm{OH}(\Delta H=13.4 \mathrm{kcal})$
b. $K_{e q}$ for the reaction $=10^{-14}$.
c. $S^{\Theta}\left(H^{\oplus}\right)=0.0$
d. $S^{\Theta}\left(\mathrm{H}_{2} \mathrm{O}\right)=16.7 \mathrm{cal} / \mathrm{molk}$.

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129. Calculate $K_{s p}$ for $A g C I$. Given:
$\Delta_{f} H^{\Theta} A g{ }^{\oplus}=25.3 \mathrm{kalmol}^{-1}$
$\Delta_{f} H^{\Theta} C 1{ }^{\Theta}=-40.0 \mathrm{kcalmol}^{-1}$
$\Delta_{f} H^{\Theta} \mathrm{AgC1}=-30.36 \mathrm{kcalmol}^{-1}$
$S^{\Theta} A g^{\oplus}, S^{\Theta} C 1{ }^{\Theta}$, and $S^{\Theta} \mathrm{AgC1are}$ 17.7, 13.2 and 23.0calmol ${ }^{-1}$

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130. Calculate the minimum mass of NaCI necessary to dissolve $0.01 \mathrm{molAgC1}$ in 100 L solution.
(Assume no
change
in
volume)
$\left(K_{f} A g C 1_{2}^{\Theta}=2.5 \times 10^{5}\right)\left(K_{s p} A G C l=2 \times 10^{-10}\right)$
131. What is the concentration of free $C d^{2+}$ in $0.005 M C d C 1_{2}$ ? $K_{1}$ for chloride complexation of $\mathrm{Cd}^{2+}$ is $100, K_{2}$ need not be considered.

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132. In the quantitative estimation of $\mathrm{Ag}^{\oplus}$ ions as AgCl , solution of NaCl is used as the precipitating reagent. Why a large excess of NaCl should be avoided?

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133. A solution was made up by $0.01 \mathrm{MCo}\left(\mathrm{NO}_{3}\right)_{2}$ and $.02 \mathrm{MN}_{2} \mathrm{H}_{4}$ and was found to have at equilibrium $\left[\mathrm{Co}^{2+}\right]=0.0062 \mathrm{M}$. Calculate $K_{1}$ for the complex formation of $\mathrm{Co}\left(\mathrm{N}_{2} \mathrm{H}_{4}\right)^{2+}$.

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134. $K_{1}, K_{2}$ and $k_{3}$ for the complexation of $\mathrm{SCN}^{\Theta}$ with $\mathrm{Fe}^{3+}$ are 130,16 , and 1.0 , respectively. (i) Calculate the overall formation costant of $\mathrm{Fe}(\mathrm{SCN})_{3}$ from its constituent ions.

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135. Explain why $0.1 \mathrm{MNH}_{3}$ solution:
a. Will precipitate $\mathrm{Fe}(\mathrm{OH})_{2}$ from 0.1 M soluton $\mathrm{Fe}^{2+}$.
b. Will not precipitate $\mathrm{Mg}(\mathrm{OH})_{2}$ from a solution which is 0.2 M in $\stackrel{\oplus}{N} \mathrm{H}_{4}$ and 0.01 M in $\mathrm{Mg}^{2+}$.
c. Will not precipitate AgOH from a solution which is 0.01 M in $\mathrm{Ag}^{\oplus}$.

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136. $K_{a}$ for $H A$ is $4.9 \times 10^{-8}$. After making the necessary approximation, calculate for its decimolar solution,

## $\Theta$

a. \% dissociation b. OH concentration
c. $p H$
137. Nicotinic acid $\left(K_{a}=1.4 \times 10^{-5}\right)$ is represented by the formula HNiC .

Calculate its percent dissociation in a solution which contains 0.10 moles of nicotinic acid per 2.0L of solution.

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138. An aqueous solution contains $10 \%$ amonia by mass and has a density of $0.99 \mathrm{gcm}^{-3}$. Calculate hydroxy1 and hydrogen ion concentration in this solution $K_{a}$ for $\mathrm{NH}_{4}^{\oplus}=5.0 \times 10^{-10} \mathrm{M}$.

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139. Determine the dergee of dissociation of $0.05 \mathrm{MNH}_{3}$ at $25^{\circ} \mathrm{C}$ in a solution of $\mathrm{pH}=11$
140. Three suggestion are made for ways to removes silver ions from solution:
a. Make the solution 0.01Min Nal.
b. Buffer the solution at $\mathrm{pH}=13$.
c. Make the solution $0.01 \mathrm{MNa}_{2} \mathrm{~S}$. What will be the equilibrium silver ion concentartion in each case? which course of action is most effective in removing $\mathrm{Ag}^{\oplus}$ ions?
$K_{\text {sp }}(\mathrm{AgI})=8.5 \times 10^{-17}, K_{\text {sp }}(\mathrm{AgOH})=2 \times 10^{-8}$,
$K_{s p}\left(A g_{2} S\right)=5.5 \times 10^{-51}$

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141. Calculate the the ratio of conjugate base// weak acid required to prepare an aqueous solution of benzoic acid and sodium benzoate with $p H$ of 4.5. The acid dissociation constant of benzoic acid is $6.5 \times 10^{-5}$.

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142. A solution is saturated with respect to $\mathrm{SrF}_{2} ; K_{s p}=7.9 \times 10^{-10}$ and $\mathrm{SrCO}_{3}, K_{\text {sp }}=7.0 \times 10^{-10}$. If the fluoride ion concentration is found to be $4.0 \times 10^{-2} \mathrm{M}$. What is the concentration of carbonates ions.

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143. The solubility of $\mathrm{Mg}(\mathrm{OH})_{2}$ in pure water is $9.57 \times 10^{-3} \mathrm{gL}{ }^{-1}$. Calculate its solubility $\mathrm{I}\left(\mathrm{gL}^{-1}\right)$ in $0.02 \mathrm{M} \mathrm{Mg}\left(\mathrm{NO}_{3}\right)_{2}$ solutions.

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144. Compare the solubility of $\mathrm{Fe}(\mathrm{OH})_{3}\left(\mathrm{~K}_{\text {sp }}=4 \times 10^{-38}\right)$ and $\mathrm{Ni}(\mathrm{OH})_{2} \cdot\left(\mathrm{~K}_{\text {sp }}=2 \times 10^{-16}\right)$ at $\mathrm{pH}=5.0$

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145. What is the $p H$ at which $0.01 M C o^{2+}$ ions in solution precipiate down as $\mathrm{Co}(\mathrm{OH})_{2} ? \mathrm{~K}_{s p}$ of $\mathrm{Co}(\mathrm{OH})_{2}$ is $2.5 \times 10^{-16}$.

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146. Freshly prepared aluminium and magnesium hydroxides are stirred vigorously in a buffer solution containing $0.25 M$ of ammonium chloride and $0.05 M$ of ammonium hydroxide. Calculate the concentration of aluminium and magesium ions in solution $\left(\mathrm{K}_{b} \mathrm{NH}_{4} \mathrm{OH}=1.8 \times 10^{-5}, K_{s p} \mathrm{Mg}(\mathrm{OH})_{2}=6 \times 10^{-10}, K_{s p} \mathrm{Al}(\mathrm{OH})_{3}=6 \times 10^{-32}\right.$

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147. A solution contains a mixture of $\mathrm{Ag}^{+}(0.10 M)$ and $\mathrm{Hg}_{2}^{2+}(0.10 M)$ which are to be separated by selective precipitation. Calculate the maximum concentration of iodide ion at which one of them gets precipitated
almost completely. $\quad\left(K_{S P} o f A g I=8.5 \times 10^{-17} \quad\right.$ and $\quad K_{S P} \quad$ of $\mathrm{Hg}_{2} \mathrm{I}_{2}=2.5 \times 10^{-26}$ )

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148. Freshly prepared aluminium and magnesium hydroxides are stirred vigorously in a buffer solution containing 0.25 M of ammonium chloride and $0.05 M$ of ammonium hydroxide. Calculate the concentration of aluminium and magesium ions in solution
$\left(\mathrm{K}_{b} \mathrm{NH}_{4} \mathrm{OH}=1.8 \times 10^{-5}, K_{s p} \mathrm{Mg}(\mathrm{OH})_{2}=6 \times 10^{-10}, K_{\text {sp }} \mathrm{Al}(\mathrm{OH})_{3}=6 \times 10^{-32}\right.$

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149. The solubility product $K_{s p}$ of $\mathrm{Ca}(\mathrm{OH})_{2}$ at $25^{\circ} \mathrm{C}$ is $4.42 \times 10^{-5} \mathrm{~A} 500$ mL of saturated solution of $\mathrm{Ca}(\mathrm{OH})_{2}$ is mixed with equal volume of 0.4 M NaOH . How much $\mathrm{Ca}(\mathrm{OH})_{2}$ in milligrams is precipitated?
150. Determine the mass of $\mathrm{PbI}_{2}$ that will dissolve in (a) 500 mL water (b) 500 mL of 0.01 MKI solution (c) 500 mL of a solution containing $1.33 \mathrm{gPb}\left(\mathrm{NO}_{3}\right)_{2}, K_{\text {sp }}$ of $\mathrm{PbI}=1.4 \times 10^{-8}$.

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151. Should a precipitate of barium fluoride be obtained when 100 mL of 0.25 MNaF and 100 mL of $0.015 \mathrm{MBa}\left(\mathrm{NO}_{3}\right)_{2}$ are mixed. $\mathrm{K}_{s p}$ of $B a F_{2}=1.7 \times 10^{-6}$

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152. A saturated solution of silver benzoate, $\mathrm{AgOCOC}_{6} \mathrm{H}_{5}$ has a pH of 8.63, $K_{a}$ for benzoic acid is $6.5 \times 10^{-5}$. Estimate $K_{s p}$ for silver benzoate.

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153. For the indicator 'HIn' the ratio $\left(\operatorname{Ind}^{\Theta}\right) /(\mathrm{HIn})$ is 7.0 at $p H$ of 4.3 . What is $K_{e q}$ for the indicator.

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154. Determine $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$ of a 0.050 M solution of ammonia to which sufficient $\mathrm{NH}_{4} \mathrm{Cl}$ has been added to make the total $\left[\mathrm{NH}_{4}^{\oplus}\right]$ equal to 0.1 M . $\left(K_{b}=1.8 \cdot 10^{-5}\right)$

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155. $K_{\text {sp }}$ of AgCl is $2.8 \times 10^{-10}$ at $25^{\circ} \mathrm{C}$. Calculate solubility of AgCl in.
a. Pure water b. $0.1 \mathrm{MAgNO}_{3}$
c. 0.1 MKCl or 0.1 MNaCl

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156. $K_{s p}$ of $\mathrm{PbC1}_{2}$ is $10^{-13}$. What will be $\left[P b^{2+}\right]$ in a of solution prepared by mixing 100 mL of $0.1 \mathrm{MPb}\left(\mathrm{NO}_{3}\right)_{2}$ of solution 1.0 mL 1 MHCI ?

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157. $K_{s p}$ of $\mathrm{PbBr}_{2}$ is $8 \times 10^{-5}$. If the salt is $80 \%$ dissociated in solution, calculate the solubility of salt in $g L^{-1}$.

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158. Equal volumes of $0.02 \mathrm{MCaC1}_{2}$ and $0.0004 \mathrm{MNa}_{2} \mathrm{SO}_{4}$ are mixed. Will a precipitate form? $K_{\text {sp }}$ for $\mathrm{CaSO}_{4}=2.4 \times 10^{-5}$ ?

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159. A solution containing both $\mathrm{Zn}^{2+}$ and $\mathrm{Mn}^{2+}$ ions at a concentration of 0.01 M is saturated with $\mathrm{H}_{2} \mathrm{~S}$. What is pH at which MnS will form a ppt ?

Under these conditions what will be the concentration of $\mathrm{Zn}^{2+}$ ions remaining in the solution ? Given $K_{s p}$ of $\mathrm{ZnSis} 10^{-22}$ and $K_{s p}$ of MnS is $5.6 \times 10^{-16}, K_{1} \times K_{2}$ of $H_{2} S=1.10 \times 10^{-21}$.

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160. For the reaction
$A g(C N)_{2}^{\ominus} \Leftrightarrow A g^{\oplus}+2 C N^{\ominus}$, the $K_{c}$ at $25^{\circ} \mathrm{C}$ is $4 \times 10^{-19}$ Calculate $\left[A g^{\oplus}\right]$ in solution which was originally 0.1 M in KCN and 0.03 M in $\mathrm{AgNO}_{3}$.

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161. A sample of hard water contains 0.05 mol of $\mathrm{CaC1}_{2}$, per litre, What is the minimum concentration of $\mathrm{Na}_{2} \mathrm{SO}_{4}$, which must be added for removing $\mathrm{Ca}^{2+}$ ions from this water sample? $K_{s p}$ for $\mathrm{CaSO}_{4}$ is $2.4 \times 10^{-5} a t 25^{\circ} \mathrm{C}$.
162. An aqueous solution of a metal bromide $\mathrm{MBr}_{2}(0.05 M)$ is saturated with $H_{2}$ S. What is the minimum pH at which MS will precipitate ? $K_{S P}$ for $M S=6.0 \times 10^{-21} \quad$ Concentration of saturqated $H_{2} S=0.1 M, K_{1}=10^{-7}$ and $K_{2}=1.3 \times 10^{-13}$ for $H_{2} S$.

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163. Calculate pH of saturated solution $\mathrm{Mg}(\mathrm{OH})_{2}, \mathrm{~K}_{\text {sp }}$ for $\mathrm{Mg}(\mathrm{OH})_{2}$ is $8.9 \times 10^{-12}$.

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164. 0.1 mili moles of $\mathrm{CdSO}_{4}$ are present in 10 ml acid solution of 0.08 NHCI . Now $\mathrm{H}_{2} \mathrm{~S}$ is passed to precipitate all the $\mathrm{Cd}^{2+}$ ions. Find the $p \mathrm{H}$ of solution after filtering off percipitate, boilling of $\mathrm{H}_{2} \mathrm{~S}$ and making the solution 100 ml by adding $\mathrm{H}_{2} \mathrm{O}$ ?

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165. The solubility of $\mathrm{Mg}(\mathrm{OH})_{2}$ is increased by the addition of $\mathrm{NH}_{4}$ ion.

Calculate
a. Kc for the reaction:
$\mathrm{Mg}(\mathrm{OH})_{2}+\stackrel{\oplus}{2 \mathrm{NH}_{4}} \Leftrightarrow 2 \mathrm{NH}_{3}+2 \mathrm{H}_{2} \mathrm{O}+\mathrm{Mg}^{+2}$
$K_{s p} o f \mathrm{Mg}(\mathrm{OH})_{2}=6 \times 10^{-12}, K_{b} o f \mathrm{NH}_{3}=1.8 \times 10^{-5}$.
b. Find the solubility of $\mathrm{Mg}(\mathrm{OH})_{2}$ in a solution containing $0.5 \mathrm{MNH}_{4} \mathrm{C} 1$ before addition of $\mathrm{Mg}(\mathrm{OH})_{2} \cdot$ b

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166. The solubility of $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ at $25^{\circ} \mathrm{C}$ is $1.20 \times 10^{-11}$. A solution of $\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ containing 0.15 mol in 500 mL water is mixed with excess of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$ till the following equilibrium is established:
$\mathrm{Ag}_{2} \mathrm{CO}_{3}+\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4} \Leftrightarrow \mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}+\mathrm{K}_{2} \mathrm{CO}_{3}$
At equilibrium, the solution constains 0.03 mol of $\mathrm{K}_{2} \mathrm{CO}_{3}$. Assuming that the degree of dissociation of $\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ and $\mathrm{K}_{2} \mathrm{CO}_{3}$ to be equal, calculate the solubility product of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$. [Take $100 \%$ ionisation of $\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ and $\left.K_{2} \mathrm{CO}_{3}\right]$

## (D) Watch Video Solution

167. 1.0 L of solution which was in equilibrium with solid mixture of $\mathrm{AgC1}$ and AgCl and $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ was found to contain $1 \times 10^{-4}$ molofAg ${ }^{\oplus}$ ions, $1.0 \times 10^{-6} \mathrm{~mol}$ of $\mathrm{C1}^{\Theta}$ ions and $8.0 \times 10^{-4}$ moles of $\mathrm{CrO}_{4}^{2-}$ ions. $\mathrm{Ag}^{\oplus}$ ions added slowely to the above mixture (keeping volume constant) till $8.0 \times 10^{-7} \mathrm{~mol}$ of $\mathrm{AgC1}$ got precipitated. How many moles of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ were also precipitated?

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168. $K_{s p}$ of $\mathrm{SrF}_{2}=2.8 \times 10^{-9}$ at $25^{\circ} \mathrm{C}$. How much NaF should be added to 100 mL of solution having 0.016 M in $\mathrm{Sr}^{2+}$ ions to reduce its concentration to $2.5 \times 10^{-3} \mathrm{M}$ ?

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169. Calculate the degree of hydrolysis and pH of 0.2 M solution of $\mathrm{NH}_{4} \mathrm{C} 1$ Given $K_{b}$ for $\mathrm{NH}_{4} \mathrm{OH}$ is $1.8 \times 10^{-5}$.

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170. Calculate for 0.01 N solution of sodium acetate,
a. Hydrolysis constant b. Dergee of hydrolysis
c. $p H$ Given $K_{a}=1.9 \times 10^{-5}$

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171. Calcium lactate is salt of weak acid and represented as $\mathrm{Ca}(\mathrm{LaC})_{2}$. A saturated solution of $\mathrm{Ca}(\mathrm{LaC})_{2}$ contains 0.13 mol of salt in 0.50 L solution. The $p O H$ of this is 5.60 . Assuming complete dissociation of salt, calculate $K_{a}$ of lacetic acid.

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172. The vapour pressur of 0.01 molal solution of weak base $B O H$ in water at $20^{\circ} \mathrm{C}$ is 17.536 mm . Calculate $K_{b}$ for base. Aqueous tension at $20^{\circ} \mathrm{C}$ is 17.540 mm . Assume molality and molarity same.

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173. Calculate the pH of $0.1 \mathrm{MK}_{3} \mathrm{PO}_{4}$ soln. The third dissociation constant of orthophosphoric acid is $1.3 \times 10^{-12}$. Assume that the hydrolysis proceeds only in the first step.

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$\oplus$
174. The ionization constant of $\mathrm{NH}_{4}$ ion in water is $5.6 \times 10^{-10}$ at $25^{\circ} \mathrm{C}$. $\oplus$ $\theta$

The rate constant the reaction of $\mathrm{NH}_{4}$ and OH ion to form $\mathrm{NH}_{3}$ and $\mathrm{H}_{2} \mathrm{O}$ at $25^{\circ} \mathrm{C}$ is $3.4 \times 10^{10} \mathrm{Lmol}^{-1} \mathrm{~S}^{-1}$. Calculate the rate constant for proton transfer from water to $\mathrm{NH}_{3}$.
175. What is $\left[\mathrm{Ag}^{+}\right]$in a solution made by dissolving both $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ and $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ until saturation is reached with respect to both salts ?
$\left[K_{\text {sp }}\left(\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}\right)=2 \times 10^{-11}, \quad K_{\text {sp }}\left(\mathrm{Ag}_{2} \mathrm{CrO}_{4}\right)=2 \times 10^{-12}\right]$

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176. Using $\mathrm{CO}_{2}, \mathrm{NH}_{3}, \mathrm{NH}_{4} \mathrm{NO}_{3}$, and $\mathrm{K}_{2} \mathrm{CrO}_{4}$ as the only reagents, devise a qualitative analysis scheme for separating and identifying the following ions, which might all be present in the same mixture: $\mathrm{Ba}^{2+}, \mathrm{Ca}^{2+}, \mathrm{Mg}^{2+}, \mathrm{Na}^{\oplus}, \mathrm{Pb}^{2+}$. Assume that each cation present is 0.10 M . State the conditions of $p H$ and the reagent concentration which are required in each step.

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177. Assuming that the only source of perodic group IIA metals is an equimolar mixture of $\mathrm{NaC1}, \mathrm{BaC1}_{2}$ and $\mathrm{MgCl}_{2}$, suggest ways of preparing
pure samples of
a. $\mathrm{MgSO}_{4}$ b. Ba metal c. $\mathrm{Ba}\left(\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}\right)_{2}$.

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178. When a solution of $\mathrm{Zn}^{2+}$ was added to a solution of NaOH , a clear solution was obtained. What $\mathrm{NH}_{4} \mathrm{C} 1$ was added to the clear solution, $\mathrm{Zn}(\mathrm{OH})_{2}$, precipitated. Using balanced chemical equation, explain these observations.

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179. Given reagents $\mathrm{NH}_{3}, \mathrm{NaOH}, \mathrm{HCl}$, and $\mathrm{H}_{2} \mathrm{~S}$ which one could be used to separated the ions in each of the following mixtures?
a. $\mathrm{Cu}^{2+}$ and $\mathrm{Zn}^{2+}$
b. $\mathrm{Cu}^{2+}$ and $A l^{3+}$
c. $\mathrm{Zn}^{2+}$ and $A l^{3+}$

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180. Estimate the $K_{\text {sp }}$ of AgBr . Given $\Delta_{f} H^{\Theta}$ of $\mathrm{Ag}^{\oplus}, \mathrm{Br}^{\Theta}$, and AgBr are 25.31, - 28.9, and -23.8kcal, $\Delta S^{\Theta}$ of $\mathrm{Ag}^{\oplus}, \mathrm{Br}^{\Theta}$, and AgBr are 17.7, 19.3, and 26.6cal/K.

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181. When 40 mL of a 0.1 M weak base, $B O H$ is titrated with 0.01 MHCl , the $p H$ of the solution at the end point is 5.5 . What will be the $p H$ if 10 mL of 0.10 MNaOH is added to the resulting solution ?

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182. Malonic acid is an organic dibasic acid such as $\mathrm{H}_{2} \mathrm{~S}$ having first ionistion constant, $K_{1}=1.42 \times 10^{-3}$ and second ionisation constant, $K_{2}=2.0 \times 10^{-6}$. Compute the divalent molanate ion concentration in:
a. 0.001 M malonic acid.
b. a solution that is $0.0001 M$ in malonic acid and $0.1 M H C 1$.
183. What mass of $p b^{2+}$ ions is left in solution, when 50.0 mL of $0.20 \mathrm{MPb}\left(\mathrm{NO}_{3}\right)_{2}$ is added to 50.0 mL of $1.5 \mathrm{MNAC1}$ ?

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184. It is given that 0.001 mol each of $\mathrm{Cd}^{2+}$ and $\mathrm{Fe}^{2+}$ ions are contained in
1.0 L of $0.02 \mathrm{MHC1}$ solution. This solutions is now saturated with $\mathrm{H}_{2} \mathrm{~S}$ gas at $25^{\circ} \mathrm{C}$.
a. Determine whether or not each of these ions will be precipitated as sulphide?
b. How much $\mathrm{Cd}^{2+}$ ions remains in the solution at equilibrium?

$$
K_{1}\left(H_{2} S\right)=1.0 \times 10^{-7}, K_{2}\left(H_{2} S\right)=1.0 \times 10^{-14}:
$$

ItbRgt
$K_{s p}(C d S)=8 \times 10^{-27}: K_{s p}(F e S)=3.7 \times 10^{-19}$.

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185. Calculate the $\left[F^{\Theta}\right]$ in a solution saturated with respect ot $M g F_{2}$ and $\mathrm{SrF}_{2}$.
$K_{\text {sp }}\left(M g F_{2}\right)=6.0 \times 10^{-9}, K_{\text {sp }}\left(S r F_{2}\right)=3.0 \times 10^{-9}$

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186. $H N_{3}$ (hydroazic acid) is a weak acid dissociating as: $H N_{3} \Leftrightarrow H^{\oplus}+N_{3}^{\Theta}$. Find the concentration of $A g^{\ominus}$ ions, if excess of solid $A g N_{3}$ is added to a solution maintained at $\mathrm{pH}=4$. The ionisation constant $K_{a}$ of $\mathrm{HN}_{3}$ is $2.0 \times 10^{-5}$. The solubility of $\mathrm{AgN}_{3}$ in pure water is found to be $5.4 \times 10^{-3}$ at $25^{\circ} \mathrm{C}$.

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187. Calculate the solubility of AgCN in a buffer solution of $p H 3.0$. Assume that no cyano complex is formed
$K_{s p} A g C N=2.2 \times 10^{-16}, K_{a} H C N=6.2 \times 10^{-10}$.
188. Find moles of $\mathrm{NH}_{4} \mathrm{Cl}$ required to prevent $\mathrm{Mg}(\mathrm{OH})_{2}$ from precipitating in a litre of solution which contains 0.02 mole $\mathrm{NH}_{3}$ and 0.001 mole $\mathrm{Mg}^{2+}$ ions.

Given : $K_{b}\left(\mathrm{NH}_{3}\right)=10^{-5}, K_{s p}\left[\mathrm{Mg}(\mathrm{OH})_{2}\right]=10^{-11}$.

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189. A solution containing $10^{-3} \mathrm{MSr}\left(\mathrm{C1O}_{4}\right)_{2}$ and $0.05 \mathrm{MKNO}_{3}$ was found to have only $75 \%$ of its strontium in the uncomplexed $\mathrm{Sr}^{2+}$ form, the rest being $\operatorname{Sr}\left(\mathrm{NO}_{3}\right)^{\oplus}$. Calcualate the $K_{1}$ for complexation reaction:
$\mathrm{Sr}^{2+}+\mathrm{NO}_{3}^{\Theta} \rightarrow \mathrm{Sr}\left(\mathrm{NO}_{3}\right)^{\Theta}$

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190. Glycine $\left[\mathrm{NH}_{2} \mathrm{CH}_{2} \mathrm{COOH}\right)$ is basic and acidic due to presence of $-\mathrm{NH}_{2}$ $\oplus$ and -COOH group. It acquires a $\mathrm{H}^{\oplus}$ to form $\mathrm{NH}_{3} \mathrm{COOH}$, which is a
diprotic acid with $K_{1}=4.55 \times 10^{-3}$ and $K_{2}=1.7 \times 10^{-10}$. In a 0.01 M solution of neutral glycine,
a. What is the pH and
b. What percent of the glycine is in the cationic form at equilibrium?

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191. A $0.01 M$ aqueous solution of weak acid $H A$ has an osotic pressure 0.293 atm at $25^{\circ} \mathrm{C}$. Another 0.01 M aqueous solution of other weak acid $H B$ has an osmotic pressure of $0.345 a t m$ under the same conditions.

Calculate equilibrium constants of two acids for their dissociation.

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192. The salt $\mathrm{Zn}(\mathrm{OH})_{2}$ is involved in the following two equilibria:
$\Theta$
$\mathrm{Zn}(\mathrm{OH})_{2}(\mathrm{~s}) \Leftrightarrow \mathrm{Zn}^{3+}(\mathrm{aq})+2 \mathrm{OH}(\mathrm{aq}) \mathrm{K}_{s p}=1.2 \times 10^{-17}$
$\mathrm{Zn}(\mathrm{OH})_{2}(\mathrm{~s})+\stackrel{\ominus}{\mathrm{O}} \mathrm{H} \Leftrightarrow \mathrm{Zn}(\mathrm{OH})_{4}^{2-}(\mathrm{aq}) \mathrm{K}_{f}=0.12$

Calculate $\left|\begin{array}{l}\Theta \\ \mathrm{OH}\end{array}\right|$ at which solubility of $\mathrm{Zn}(\mathrm{OH})_{2}$ be a minimum. Also find the solubility of $\mathrm{Zn}(\mathrm{OH})_{2}$ at this pH .

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193. A 500 mL of an equilibrium mixture of gaseous $\mathrm{N}_{2} \mathrm{O}_{4}$ and $\mathrm{NO}_{2}$ at $25^{\circ} \mathrm{C}$ and 753 mm of Hg was allowed to react with enough water to make 250 mL of solution at $25^{\circ} \mathrm{C}$. Assume that all the dissolved $\mathrm{N}_{2} \mathrm{O}_{4}$ is converted to $\mathrm{NO}_{2}$ which disproportionates in water yielding a solution of nitrous acid and nitric acid. aAsume further that disproportionation reaction goes to completion and that none of the nitrous acid disproportionates. The equilibrium constant $\left(K_{p}\right)$ for $\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \Leftrightarrow 2 \mathrm{NO}_{2}(\mathrm{~g}) 0.113$ at $25^{\circ} \mathrm{C} . \mathrm{K}_{a}$ for $\mathrm{HNO}_{2}$ is $4.5 \times 10^{-4}$ at $25^{\circ} \mathrm{C}$.
a. Write balanced equation for disproportionation.
b. What is the molar concentration of $\mathrm{NO}_{2}$ and pH of the solution?
c. What is osmotic pressure of solution?
d. How many grams of lime ( CaO ) would be required to neutralise the solution?
194. A buffer solution 0.04 M in $\mathrm{Na}_{2} \mathrm{HPO}_{4}$ and 0.02 in $\mathrm{Na}_{3} \mathrm{PO}_{4}$ is prepared. The electrolytic oxidation of 1.0 milli-mole of the organic compound RNHOH is carried out in 100 mL of the buffer. The reaction is $\mathrm{RNHOH}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{RNO}_{2}+4 \mathrm{H}^{+}+4 e^{-}$The approximate pH of solution after the oxidation is complete is :
[Given: $f$ or $H_{3} \mathrm{PO}_{4}, p K_{a 1}=2.2, p K_{a 2}=7.20, p K_{a 3}=12$ ]
(a) 6.90
(b) 7.20
(c) 7.5
(d)None of these

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195. Calculate the pH of $0.05 \mathrm{MKHC}_{8} \mathrm{H}_{4} \mathrm{O}_{4}$

$$
\begin{aligned}
& \mathrm{H}_{2} \mathrm{C}_{8} \mathrm{H}_{4} \mathrm{O}_{4}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{H}_{3} \mathrm{O}^{\oplus}+\mathrm{HC}_{8} \mathrm{H}_{4} \mathrm{O}_{4}^{\oplus} p K_{a_{1}}=2.94 \\
& \mathrm{HC}_{8} \mathrm{H}_{4} \mathrm{O}_{4}^{\Theta}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{H}_{3} \mathrm{O}^{\oplus}+\mathrm{C}_{8} \mathrm{H}_{4} \mathrm{O}_{4}^{2-} p K_{a_{2}}=5.44
\end{aligned}
$$

196. It is found that 0.1 M solution of three sodium salts $\mathrm{NaX}, \mathrm{NaY}$, and NaZ have pH 7.0, 9.0 and 11.0, respectively. Arrange the acids (HX, HY, and HZ ) in order of increasing acidic character. Calculate dissociation constant of acids.

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197. What is $\left[\mathrm{Cd}^{2+}\right]$ in 1.0 L of solution prepared by dissolving $0.001 \mathrm{molCd}\left(\mathrm{NO}_{3}\right)_{2}$ and 1.5mmol. $\mathrm{NH}_{3}$ ? $\mathrm{K}_{d}$ for the dissociation of $\mathrm{Cd}\left(\mathrm{NH}_{3}\right)_{4}^{2+}$ into $\mathrm{Cd}^{2+}$ and $4 \mathrm{NH}_{3}$ is $1.8 \times 10^{-7}$. Neglect the amount of Cd in complexes containing fewer than 4 ammonia molecules.

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198. 0.001 mol of solid $\mathrm{NaC1}$ was added to 1.0 L of $0.01 \mathrm{MHg}\left(\mathrm{NO}_{3}\right)_{2}$. Calculate $\left[c 1^{\Theta}\right]$ equilibrated with newly formed $\mathrm{HgC1}{ }^{\oplus} . \mathrm{K}_{1}$ for $\mathrm{HgC1}{ }^{\oplus}$ formation is $5.5 \times 10^{6}$, neglect the $K_{2}$ equilibrium.

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199. How much $\mathrm{NH}_{3}$ should be added to a solution of $0.01 \mathrm{MCu}\left(\mathrm{NO}_{3}\right)_{2}$ to reduce $\left[\mathrm{Cu}^{2+}\right.$ to $10^{-13}$. Neglect the amount of copper in complexes containing fewer than 4 ammonia molecules per copper atom. Given $K_{d}$ for $\mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{4}^{2+}=1.0 \times 10^{-12}$

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200. Calculated the minimum amount of $N h_{3}$ which must be added to
1.0 L of solution in order to dissolve $0.1 \mathrm{molAgC1}$ by forming $\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{\oplus} ? \mathrm{~K}_{\text {sp }}$ of $\mathrm{AgC1}=1 \times 10^{-10}, K_{f}\left(\mathrm{NH}_{3}\right)_{2}^{\oplus}=1 \times 10^{8}$.
201. A certain insoluble compound of $M^{2+}$, when shaken with water, provides an $\mathrm{M}^{2+}$ concentration of $1.0 \times 10^{-4} \mathrm{M}$. A ligand is added to the system in a quantify which forms a soluble complex with $\mathrm{M}^{2+}$ and leaves $1.0 \times 10^{-6} \mathrm{M}, \mathrm{M}^{2+}$ in solution. Will the insoluble compound tend to dissolve? Explain.

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## Ex 8.1

1. 100 mL of HCl gas at $25^{\circ} \mathrm{C}$ and 740 mm pressure is dissolved in 1 L of $\mathrm{H}_{2} \mathrm{O}$. Calculate the pH of solution. Given vapour presure of $\mathrm{H}_{2} \mathrm{O}$ at $25^{\circ} \mathrm{C}$ is 23.7 mm .

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2. Calculate $\left[C l^{\Theta}\right],\left[N a^{\oplus}\right],\left[H^{\oplus}\right],\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$, and the $p H$ of resulting solution obtained by mixing 50 mL of 0.6 MHCl and 50 mL of 0.3 MNaOH .

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3. Calculate the pH of solution obtained by mixing 10 ml of 0.1 MHCl and 40 ml of $0.2 \mathrm{MH}_{2} \mathrm{SO}_{4}$

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4. Calculate the pH of a solution which contains 100 mL of $0.1 \mathrm{MHC1}$ and 9.9mL of 1.0 MNaOH .

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5. Calculate the $\left[H^{\oplus}\right]$ and $\left[\begin{array}{l}\Theta \\ \mathrm{OH}\end{array}\right]$ of 0.0315 g of $\mathrm{HNO}_{3}$ in 500 mL of water.

Calculate pH and pOH also.

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6. A 25.0 mL . sample of 0.10 M HCl is titrated with 0.10 M NaOH . What is the pH of the solution at the points where 24.9 and 25.1 mL of NaOH have been added?

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7. Conjugate acid of $\mathrm{NH}_{2}$ :
A. $\mathrm{N}_{2} \mathrm{H}_{4}$
B. $\mathrm{NH}_{3}$
C. $\mathrm{NH}_{2} \mathrm{OH}$
$\Theta$
D. $\mathrm{NH}_{4}$

## Answer: B

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8. Which of the following solutions will have pH close to 1.0 ?
A. 75 mL of $\mathrm{M} / 5 \mathrm{HC} 1+25 \mathrm{~mL}$ of $\mathrm{M} / 5 \mathrm{NaOh}$
B. 10 mL of $\mathrm{M} / 10 \mathrm{HC} 1+90 \mathrm{~mL}$ of $\mathrm{M} / 10 \mathrm{NaOH}$
C. 55 mL of $\mathrm{M} / 10 \mathrm{HC} 1+54 \mathrm{~mL}$ of $\mathrm{M} / 10 \mathrm{NaOH}$
D. 100 mL of $\mathrm{M} / 10 \mathrm{HC} 1+100 \mathrm{~mL}$ of $\mathrm{M} / 10 \mathrm{NaOH}$

## Answer: A

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9. An acid solution of $p H=6$ is diluted 100 times. The $p H$ of solution
A. (a) 6.95
B. (b) 6
C. (c) 4
D. (d) 8

## Answer: A

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10. The number of $H^{\oplus}$ ions present in 1 mL of solution having $p H=13$ is
A. (a) $6.023 \times 10^{10}$
B. (b) $6.023 \times 10^{7}$
C. (c) $6.023 \times 10^{13}$
D. (d) $10^{13}$

## Answer: B

11. Equal volumes of two solutions of HCl are mixed. One solution has a $p H=1$, while the other has a $p H=5$. The $p H$ of the resulting solution is
A. $<1$
B. Between 1 and 2
C. 3
D. Between 4 and 5

## Answer: B

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12. For pure water:
A. Both pH and pOH decrease with increase in temperature.
B. Both pH and pOH increase with increase in temperature.
C. $p H$ decreases and $p O H$ increases with increase in temperature.
D. pH increase and pOH decreases with increase in temperature.

## Answer: A

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13. The $p H$ of a solution increased from 3 to 6 . Its $\left[H^{\oplus}\right]$ will be
A. Reduced by 1000 times
B. Increased to 1000 times
C. Doubled
D. Reduced to half

## Answer: A

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14. The following equilibria is established when $\mathrm{H}_{2} \mathrm{SO}_{4}$ is dissolved in acetic acid:
$\mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{CH}_{3} \mathrm{COOH} \Leftrightarrow \stackrel{\ominus}{\mathrm{H}} \mathrm{O}_{4}+\mathrm{CH}_{3} \mathrm{COOO}_{2}{ }_{2}$
The set that characterised the conjugate acid-base pairs is:
A. $\left(\mathrm{H}_{2} \mathrm{SO}_{4}, \mathrm{CH}_{3} \mathrm{COOH}\right)$ and $\left(\mathrm{CH}_{3} \mathrm{COOH}_{2}, \mathrm{HSO}_{4}\right)$
B. $\left(\mathrm{H}_{2} \mathrm{SO}_{4}, \mathrm{CH}_{3} \mathrm{COO} \stackrel{\oplus}{\mathrm{H}_{2}}\right)$ and $\left(\mathrm{CH}_{3} \mathrm{COOH}, \mathrm{H}_{2} \mathrm{SO}_{4}\right)$
c. $\left(\mathrm{CH}_{3} \mathrm{COOH}_{2}{ }^{\oplus}, \mathrm{H}_{2} \mathrm{SO}_{4}\right)$ and $\left(\stackrel{\ominus}{\mathrm{HsO}_{4}}, \mathrm{CH}_{3} \mathrm{COOH}\right)$
D. $\left(\mathrm{H}_{2} \mathrm{SO}_{4}, \stackrel{\ominus}{\mathrm{HSO}_{4}}\right)$ and $\left(\mathrm{CH}_{3} \mathrm{COOH}_{2}, \mathrm{CH}_{3} \mathrm{COOH}\right)$

## Answer: C

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15. Which is the set of amphiprotic species?
A. $\mathrm{H}_{2} \mathrm{O}, \mathrm{H}_{2} \mathrm{PO}_{3}^{\Theta}, \mathrm{HPO}_{4}^{2-}$
B. $\mathrm{HC}_{2} \mathrm{O}_{4}^{\Theta}, \mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}, \mathrm{SO}_{4}^{2-}$
C. $\mathrm{H}_{2} \mathrm{O}, \mathrm{HPO}_{4}^{2-}, \mathrm{H}_{2} \mathrm{PO}_{2}^{\Theta}$
D. $\mathrm{H}_{3} \mathrm{O}^{\oplus}, \mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}, \mathrm{HCO}_{3}^{\Theta}$

## Answer: A

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16. One litre of 0.5 MKCl is electrolysed by passing 9650 coulombs of electricity. The ${ }^{`} \mathrm{pH}$ of resulting solution is
A. 1.0
B. 2.0
C. 7.0
D. 13.0
17. pH of a solution made by mixing 200 mL of $0.0657 \mathrm{MNaOH}, 140 \mathrm{~mL}$ of
0.107 MHCl and 160 mL of $\mathrm{H}_{2} \mathrm{O}$ is
A. 3.04
B. 2.43
C. 2.74
D. 2.27

## Answer: B

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18. When one drop of a concentrated HCl is added to 1 L of pure water at $25^{\circ} \mathrm{C}$, the pH drops suddenly from 7 to 4 . When the second drop of the same acid is added, the $p H$ of the solution further drops to about
A. 0
B. 1.0
C. 2.0
D. 3.7

## Answer: D

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## Ex 8.2

1. The dissociation constant of acetic acid is $8 \times 10^{-5}$ ta $25^{\circ} \mathrm{C}$. Find the pH of
i. $M / 10 \mathrm{ii} . M / 100$ solution of acetic acid.
2. Calculate the amount of acetic acid present in $1 L$ of solution having $\alpha=1 \%$ and $K_{a}=1.8 \times 10^{-5}$.

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3. 0.16 g of $\mathrm{N}_{2} \mathrm{H}_{4}$ are dissolved in water and the total volume made upto 500 mL . Calculate the percentage of $\mathrm{N}_{2} \mathrm{H}_{4}$ that has reacted with water in this solution. $\left(K_{b}\right.$ for $\left.N_{2} H_{4}=4.0 \times 10^{-6}\right)$

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4. If the pH of $0.26 \mathrm{MHNO}_{2}$ is 2.5 , what will be its dissociation constant.

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5. Find the dissociation constant $K_{a}$ of $H A$ (weak monobasic acid) which is $3.5 \%$ dissociated in an $M / 20$ solution.
6. Ionic product of water $\left(K_{w}\right.$ is $\left.10^{-14}\right)$ at $25^{\circ} \mathrm{C}$. What is the dissociation constant of water and auto protonation constant of water?

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7. 2.0 gof diborane $\left(\mathrm{B}_{2} \mathrm{H}_{6}\right)$ reacts with water to product 100 mL solution. If $K_{a}$ for $\mathrm{H}_{3} \mathrm{BO}_{3}$ is $7.3 \times 10^{-10}$, calculated the pH of solution.

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8. At $90^{\circ} \mathrm{C}$, pure water has $\left[\mathrm{H}_{3} \mathrm{O}^{\oplus}\right]=10^{-6} \mathrm{M}$. What is the value of $K_{w}$ at $90^{\circ} \mathrm{C}$
A. $10^{-6}$
B. $10^{-8}$
C. $10^{-12}$
D. $10^{-14}$

## Answer: C

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9. HCOOH and $\mathrm{CH}_{3} \mathrm{COOH}$ solutions have equal pH . If $K_{1} / K_{2}$ is 4 , the ratio of their molar concentration will be
A. 0.25
B. 0.5
C. 2
D. 4

## Answer: A

10. $2 \mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{H}_{3} \mathrm{O}^{\oplus}+\mathrm{OH}, K_{w}=10^{-14}$ at $25^{\circ} \mathrm{C}$, hence $K_{a}$ is
A. $10^{-7}$
B. $5.55 \times 10^{-13}$
C. $10^{-14}$
D. $18 \times 10^{-17}$

## Answer: D

## D Watch Video Solution

11. Which of the following expression is wrong?
A. $\left[H^{\oplus}\right]=\left[\begin{array}{l}\Theta \\ O H\end{array}\right]=10^{-7}$ for a neutral solution at all temperatures.
B. $\left[H^{\oplus}\right]<\sqrt{K_{w}}$ and $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]>\sqrt{K_{w}}$ for basic solution
C. $\left[H^{\oplus}\right]=\left[\begin{array}{l}\Theta \\ O H\end{array}\right]=\sqrt{K_{w}}$ for a neutral solution
D. $\left[H^{\oplus}\right]>\sqrt{K_{w}}$ and $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]<\sqrt{K_{w}}$ for an acidic solution

## Answer: A

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12. For a ' $C$ ' $M$ concentrated solution of a weak electrolyte ' $A_{-}(x) B_{-}(y)$ 'alpha' degree of dissociation) is
A. $\sqrt{K_{e q} / x y C}$
B. $\left[K_{e q} \cdot C /(x y)\right]^{1 / 2}$
C. $\left(K_{e q} / C^{x+y-1} \cdot x^{x} x^{y}\right)^{\frac{1}{x+y}}$
D. $\left[K_{e q} / C(x+y)\right]^{1 / 2}$

## Answer: C

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13. $\mathrm{K}_{b}$ for $\mathrm{NH}_{4} \mathrm{OH}$ is $1.8 \times 10^{-5}$. The $\left[\begin{array}{l}\Theta \\ \mathrm{OH}\end{array}\right]$ of $0.1 \mathrm{MNH}_{4} \mathrm{OH}$ is
A. $5.0 \times 10^{-2}$
B. $4.20 \times 10^{-3}$
C. $1.34 \times 10^{-3}$
D. $1.8 \times 10^{-6}$

## Answer: C

## D Watch Video Solution

14. The dissociation constant of monobasic acids $A, B, C$ and $D$ are $6 \times 10^{-4}, 5 \times 10^{-5}, 3.6 \times 10^{-6}$, and $7 \times 10^{-10}$, respectively. The $p H$ values of their 0.1 M aqueous solutions are in the order.
A. $D>C>B>A$
B. $A>B>C>D$
C. $A=B=C=D$
D. None

## Answer: A

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15. The molarity of $\mathrm{NH}_{3} \mathrm{OfpH}=12 \mathrm{at} 25^{\circ} \mathrm{C}$ is $\left(K_{b}=1.8 \times 10^{-5}\right)$
A. 11.7 M
B. 5.5 M
C. $0.55 M$
D. 0.01 M

## Answer: C

16. A weak acid, HA, has a $K_{a}$ of $1.00 \times 10^{-5}$. If 0.100 mol of the acid is dissolved in 1 L of water, the percentage of the acid dissociated at equilibrium is the close to
A. $0.1 \%$
B. $1.0 \%$
C. 99.0 \%
D. 99.9 \%

## Answer: B

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17. For a polyprotic acid, $\mathrm{H}_{3} \mathrm{PO}_{4}$ its three dissociation constanst $K_{1}, K_{2}$ and $K_{3}$ are in the order
A. $K_{1}=K_{2}>K_{3}$
B. $K_{1}=K_{2}=K_{3}$
C. $K_{1}>K_{2}>K_{3}$
D. $K_{1}<K_{2}<K_{3}$

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18. Given $\mathrm{HF}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{F}^{-}: \mathrm{K}_{a}$

$$
\mathrm{F}^{-}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{HF}+\mathrm{OH}^{-}, K_{b}
$$

Which of the following relations is correct ?
A. $K_{b}=K_{w}$
B. $K_{a} \times K_{b}=K_{w}$
C. $K_{b}=\frac{1}{K_{w}}$
D. $\frac{K_{a}}{K_{b}}=K_{w}$

## Answer: B

19. A certain weak acid has a dissociation constant $1.0 \times 10^{-4}$. The equilibrium constant for its reaction with a strong base is :
A. $10^{-14}$
B. $10^{-9}$
C. $10^{-10}$
D. $10^{10}$

## Answer: D

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20. The percentage error in $\left[H^{\oplus}\right]$ provided by $10^{-8} \mathrm{MHC1}$, if ionisation of water is not neglected, is
A. $2 \%$
B. $3 \%$
C. $4 \%$
D. $5 \%$

## Answer: D

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21. $\mathrm{H}_{3} \mathrm{BO}_{3}$ is :
A. Monobasic and weak Lewis acid
B. Monobasic and weak Bronsted acid
C. Tribasic and weak Bronsted acid
D. Monobasic and strong Lewis acid

## Answer: A

## - Watch Video Solution

22. The enthalpy change for first proton neutralisation of $\mathrm{H}_{2} \mathrm{~S}$ is $-37.1 \mathrm{kJmol}^{-1}$. What is the enthalpy change for first ionisation of ${ }^{\mathrm{H}} \mathrm{H}_{-}(2) \mathrm{S}$.' in $\mathrm{KJ} / \mathrm{mol}$.
A. 94.2
B. -20.0
C. 20.0
D. -94.2

## Answer: C

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23. The factor by which the degree of ionisation of 200 mL of 0.1 M benzoic acid solution $\left(K_{a}=4 \times 10^{-5}\right)$ changes on addition of 100 mL of 0.2 MHCl is:
A. 0.02
B. 0.03
C. 33.33
D. None

## Answer: B

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24. The concentration of $\mathrm{CO}_{2}$ in atmosphere is 88 ppm . If all of the $\mathrm{CO}_{2}$ present in $10^{5} \mathrm{~mL}$ of air is dissolved in $1 \mathrm{dm}^{3}$ water, then approximate pOH of solution at $27^{\circ} \mathrm{C}$ will be $\left(K_{a_{1}}=10^{-7}, K_{a_{2}}=10^{-11}\right.$ for $\left.\mathrm{H}_{2} \mathrm{CO}_{3}\right]$
A. 3.2
B. 3.85
C. 10.15
D. None
25. A solution of a weak monoprotic acid has dissociation constant $K_{a}$. The minimum initial concentration $C$ such that the concentration of the undissociated acid can be equated to $C$ within an error of $1 \%$ would be
A. $9900 K_{a}$
B. $10000 K_{a}$
C. $99 K_{a}$
D. $K_{a}$

## Answer: A

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26. Two weak acids $H X$ and $H Y$ have $K_{a}$ values $1.75 \times 10^{-5}$ and $1.3 \times 10^{-5}$, respectively, at a certain temperature. An equimolar solution of mixture
of two acids is parially neutralised by NaOH . How is the ratio of the contents of $X^{\Theta}$ and $Y^{\Theta}$ ions related to the $K_{a}$ values and molarity?
A. $\left[\frac{\alpha}{1-\alpha}\right]=\frac{1.75}{1.3} \times\left[\frac{\alpha^{\prime}}{1-\alpha^{\prime}}\right], 0$, where $\alpha$ and $\alpha^{\prime}$ are ionised fractions of the acids $H X$ and $H Y$ respectively.
B. The ratio is unrelated to the $K_{a}$ values.
C. The ratio is unrelated to the molarity.
D. The ratio is unrelated to the pH of the solution.

## Answer: A

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27. $\mathrm{H}_{2} \mathrm{~S}$ behaves as a weak diprotic acid in aqueous solution. Which of the following is the correct explanantion for pH of a solution of $\mathrm{H}_{2} \mathrm{~S}$ in terms of its $p K_{1}, p K_{2},\left[H_{2} S\right]$ and $\left[S^{2-}\right]$
A. $p H=\frac{1}{2}\left(p K_{1}+p K_{2}\right)$
B. $p H=\frac{1}{2}\left(p K_{1}+p K_{2}-\log \frac{\left[S^{2-}\right]}{\left[H_{2} S\right]}\right)$
C. $p H=\frac{1}{2}\left(p K_{1}+p K_{2}+\log \frac{\left[S^{2-}\right]}{\left[H_{2} S\right]}\right)$
D. $p H=\frac{1}{2}\left(p K_{1}-p K_{2}+\log \frac{\left[\mathrm{H}_{2-} \mathrm{S}\right]}{\left[S^{2-}\right]}\right)$

## Answer: C

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## Ex 8.3

1. Calculate the pH of a solution of given mixture.
$\left(2 g \mathrm{CH}_{3} \mathrm{COOH}+3 g \mathrm{CH}_{3} \mathrm{COONa}\right)$ in 100 mL of mixture.

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2. How many moles of $\mathrm{NH}_{4} \mathrm{Cl}$ should be added to 200 mL solution of $0.18 \mathrm{MNH}_{4} \mathrm{OH}$ to have a pH of $9.60 . \mathrm{K}_{b}$ of $\mathrm{NH}_{4} \mathrm{OH}=2 \times 10^{-5}$

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3. A buffer solution was made by adding 15.0 g of $\mathrm{CH}_{3} \mathrm{COOH}$ and $20.5 \mathrm{gCH}_{3} \mathrm{COONa}$. The buffer is diluted to 1.0 L .

Calculate the pH of solution.
Given: $p K_{a} o f \mathrm{CH}_{3} \mathrm{COOH}=4.74, \log \left(\frac{13}{12}\right)=0.035$

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4. A buffer solution contains $0.25 \mathrm{MNH}_{4} \mathrm{OH}$ and $0.3 \mathrm{NH}_{4} \mathrm{C} 1$.
a. Calculate the pH of the solution.
$K_{b}=2 \times 10^{-5}$.
5. Calculate the hydrolysis constant $\left(K_{h}\right)$ and degree of hydrloysis (h) of $\mathrm{NH}_{4} \mathrm{C} 1$ in 0.1 M solution.
$K_{b}=2.0 \times 10^{-5}$. Calculate the $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$ ions in the solution.

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6. Calcualte the percentage hydrolysis of $10^{-3} \mathrm{MN}_{2}^{\oplus} \mathrm{H}_{5} \mathrm{C1}{ }^{\Theta}$ (hydrazinium chloride), salt contining acid ion conjugate to hydrazine base $\left(\mathrm{NH}_{2} \mathrm{NH}_{2}\right) \cdot K_{b}$ for $\mathrm{N}_{2} \mathrm{H}_{4}=1.0 \times 10^{-6}$.

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7. Calculate the amount of $\mathrm{NH}_{4} \mathrm{Cl}$ required to dissolve in 500 mL of water to have a $\mathrm{pH}=4.5, K_{b}=2.0 \times 10^{-5}$.

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8. A 0.25 M solution of pyridinium chloride $\left(\mathrm{C}_{5} \mathrm{H}_{5}{ }^{\oplus} \mathrm{HCl} \mathrm{H}^{\Theta}\right)$ has pH of 2.89 .

Calculate $p K_{b}$ for pyridine $\left(\mathrm{C}_{5} \mathrm{H}_{5} \mathrm{~N}\right)$.

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9. Which of the following is a buffer solution?
A. A solution of $\mathrm{KC1}$ and KOH
B. A solution of $\mathrm{CH}_{3} \mathrm{COONH}_{4}$
C. A solution of $\mathrm{K}_{2} \mathrm{SO}_{4}$ and $\mathrm{NH}_{4} \mathrm{OH}$
D. A solution of PhCOOK nad PhCOOH

## Answer: D

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10. Which of the following is not a buffer?
A. $\mathrm{NH}_{4} \mathrm{OH}+\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOH}$
B. $\mathrm{HCOOH}+\mathrm{HCOONa}$
C. $\mathrm{H}_{2} \mathrm{CO}_{3}+\mathrm{HCOOK}$
D. $\mathrm{NH}_{4} \mathrm{OH}+\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4}$

## Answer: A

## D Watch Video Solution

11. In an acidic buffer solution, if some $\mathrm{H}_{2} \mathrm{So}_{4}$ is added, its pH will
A. Remain constant
B. Change but cannot be predicted
C. Decrease
D. Increase

## Answer: C

12. Which of the following solutions containing weak acid and salt of its conjugate base has maximum buffer capaity?
A. [Salt] $<$ [Acid]
B. $[$ Salt $]=[$ Acid $]$
C. [Salt] > [Acid]
D. [Salt] + [Acid] is minimum

## Answer: B

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13. A weak acid $H A$ has $K_{a}=10^{-6}$. What would be the molar ratio of this acid and its salt with strong base so that $p H$ of the buffer solution is 5 ?
A. $1 / 10$
B. 10
C. 1
D. 2

## Answer: B

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14. The addition of $\mathrm{NaH}_{2} \mathrm{PO}_{4}$ to $0.1 \mathrm{MH}_{3} \mathrm{PO}_{4}$ will cuase
A. No change in $p H$ value
B. Increases in its pH value
C. Decrease in its pH value
D. Change in $p H$ but cannot be predicted

## Answer: B

## - Watch Video Solution

15. On diluting a buffer solution, its pH
A. Increases
B. Decreases
C. Remains same
D. Cannot be predicted

## Answer: C

## - Watch Video Solution

16. The pH of a solution containing 0.1 mol of $\mathrm{CH}_{3} \mathrm{COOH}, 0.2 \mathrm{~mol}$ of $\mathrm{CH}_{3} \mathrm{COONa}$, and 0.05 mol of NaOH in $1 \mathrm{~L} .\left(p \mathrm{~K}_{a} o f \mathrm{CH}_{3} \mathrm{COOH}=4.74\right)$ is:
A. 5.44
B. 5.20
C. 5.04
D. 4.74

## Answer: A

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17. A weak base $B O H$ is titrated with strong acid $H A$. When $10 m L$ of $H A$ is added, the $p H$ is 9.0 and when $25 m L$ is added, $p H$ is 8.0 . The volume of acid required to reach the equivalence point is
A. 50 mL
B. 40 mL
C. $35 m L$
D. 30 mL

## Answer: D

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18. To 1.0 L solution containing 0.1 mol each of $\mathrm{NH}_{3}$ and $\mathrm{NH}_{4} \mathrm{C} 1,0.05 \mathrm{molNaOH}$ is added. The change in $p H$ will be $\left(p K_{a}\right.$ for $\mathrm{CH}_{3} \mathrm{COOH}=4.74$ )
A. 0.30
B. -0.30
C. 0.48
D. -0.48

## Answer: C

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19. The pH of blood is 7,4 . If the buffer in blood constitute $\mathrm{CO}_{2}$ and $\mathrm{HCO}_{3}^{\Theta}$ ions, calculate the ratio of conjugate base of acid $\left(\mathrm{H}_{2} \mathrm{CO}_{3}\right)$ to maintain the pH of blood. Given $K_{1}$ of $\mathrm{H}_{2} \mathrm{CO}_{3}=4.5 \times 10^{-7}$.
A. 11.25
B. 10.0
C. 8.5
D. None

## Answer: A

## - Watch Video Solution

20. The pH of blood is
A. $>10$
B. Between 8 and 0
C. Between 7 and 8
D. $<6$

## Answer: C

21. Buffer in blood consists of
A. $\mathrm{H}_{2} \mathrm{CO}_{3}$ and $\mathrm{Cl}^{-}$
B. HCl and $\mathrm{HCO}_{3}^{-}$
C. HCl and $\mathrm{Cl}^{-}$
D. $\mathrm{H}_{2} \mathrm{CO}_{3}$ and $\mathrm{HCO}_{3}^{-}$

## Answer: D

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22. $K_{a}$ for $H C N$ is $5 \times 10^{-10}$ at $25^{\circ} \mathrm{C}$. For maintaining a constant $p H$ of 9.0 , the volume of $5 M K C N$ solution required to be added to 10 mL of $2 M H C N$ solution is
A. $9.3 m L$
B. 7.95 mL
C. $4 m L$
D. $2 m L$

## Answer: D

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23. 18 mL of mixture of $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COONa}$ required 6 mL of 0.1 MNaOH for neutralisation of the acid 12 mL of 0.1 MHCl for reaction with salt, separately. If $p K_{a}$ of the acid is 4.75 , what is the $p H$ of the mixture
A. 4.5
B. 4.6
C. 4.75
D. 5.05

## Answer: D

24. The pH of blood is maintained by the balance between $\mathrm{H}_{2} \mathrm{CO}_{3}$ and $\mathrm{NaHCO}_{3}$. If the amount of $\mathrm{CO}_{2}$ in the blood is increased, how will it effect the pH of blood'?
A. $p H$ will remain same.
B. $p H$ wil be 7
C. $p H$ will increases.
D. $p H$ will decrease.

## Answer: A

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25. Fixed volume of 0.1 M benzoic acid $\left(p K_{a}=4.2\right)$ solution is added into 0.2 M sodium benzote solution and formed a 300 mL , resulting acidic buffer solution. If $p H$ of the resulting solution is 3.9 , then added volume of banzoic acid is
A. $240 m L$
B. 150 mL
C. 100 mL
D. None

## Answer: A

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26. 0.1 mol of $\mathrm{RNH}_{2}\left(K_{b}=5 \times 10^{-4}\right)$ is mixed with 0.08 mol of $\mathrm{HC1}$ and diluted to 1 L . Calculate the $\left[H^{\oplus}\right]$ in the solution.
A. $8 \times 10^{-11} M$
B. $1.6 \times 10^{-11} M$
C. $8 \times 10^{-5} M$
D. $8 \times 10^{-2} M$

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27. A weak acid $H X\left(K_{a}=10^{-5}\right)$ on reaction with NaOH gives NaX . For 0.1 M aqueous solution of NaX , the \% hydrolysis is
A. $1 \%$
B. 0.01 \%
C. 0.001 \%
D. $0.15 \%$

## Answer: B

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28. The pH of 0.1 M solution of the following salts decreases in the order
A. $\mathrm{HCl}>\mathrm{NaCl}>\mathrm{NH}_{4} \mathrm{Cl}>\mathrm{NaCN}$
B. $\mathrm{HCl}>\mathrm{NaCN}>\mathrm{NH}_{4} \mathrm{Cl}>\mathrm{NaCl}$
C. $\mathrm{NaCN}>\mathrm{NaCl}>\mathrm{NH}_{4} \mathrm{Cl}>\mathrm{HCl}$
D. $\mathrm{NH}_{4} \mathrm{Cl}>\mathrm{NaCN}>\mathrm{NaCl}>\mathrm{HCl}$

## Answer: C

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29. The degree of hydrolysis of a salt of $W_{A}$ and $W_{B}$ in its $0.1 M$ solution is $50 \%$. If the molarity of the solution is 0.2 M , the percentage hydrolysis of the salt woukd be
A. $25 \%$
B. 50 \%
C. 75 \%
D. $100 \%$

## Answer: B

30. $p H$ of separate solution of four potassium salts, $K W, K X, K Y$ and $K Z$ are $7.0,9.0,10.0$, and 10.5 , respectively. If each solution is $0.2 M$, the strongest acid would be
A. $H W$
B. $H X$
C. $H Y$
D. $H Z$

## Answer: A

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31. Which of the following solutions have $\mathrm{pH}<7$.
A. $\mathrm{BaI}_{2}$
B. $\mathrm{AI}\left(\mathrm{NO}_{3}\right)_{3}$
C. $\mathrm{CH}_{3} \mathrm{COONH}_{4}$
D. CsI

## Answer: B

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32. Which of the following solution have $\mathrm{pH}>7$.
I. $B a F_{2}$ II. $R b I$
III. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COONa}$
IV. $\left(\mathrm{CH}_{3} \mathrm{COO}\right)_{2} \mathrm{Ba}$
A. I
B. I, II, III
C. I, III
D. I, III, IV

## Answer: D

33. The expression to calculate pH of sodium acetate solution at $25^{\circ} \mathrm{C}$ is
A. $p \mathrm{H}=7+\frac{1}{2} p K_{b}\left(\mathrm{CH}_{3} \mathrm{COOH}\right)-\frac{1}{2} \log [$ salt $]$
B. $\mathrm{pH}=7+\frac{1}{2} p K_{a}\left(\mathrm{CH}_{3} \mathrm{COOH}\right)-\frac{1}{2} \log [$ salt $]$
C. $\mathrm{pH}=7+\frac{1}{2} p K_{b}\left(\mathrm{CH}_{3} \mathrm{COOH}\right)+\frac{1}{2} \log [$ salt $]$
D. $\mathrm{pH}=7+\frac{1}{2} p K_{a}\left(\mathrm{CH}_{3} \mathrm{COOH}\right)+\frac{1}{2} \log [$ salt $]$

## Answer: D

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34. The correct order of increasing $\left[\mathrm{H}_{3} \mathrm{O}^{\oplus}\right]$ in the following aqueous solution is
A. $0.01 \mathrm{MH}_{2} \mathrm{~S}<0.01 \mathrm{MH}_{2} \mathrm{SO}_{4}<0.01 \mathrm{MNaC1}<0.01 \mathrm{MNaNO}_{3}$
B. $0.01 \mathrm{MNaC1}=0.01 \mathrm{MNaNO}_{3}<0.01 \mathrm{MH}_{2} \mathrm{~S}<0.01 \mathrm{MH}_{2} \mathrm{SO}_{4}$
C. $0.01 \mathrm{MH}_{2} \mathrm{~S}<0.01 \mathrm{MNaNO}_{3}=0.01 \mathrm{MNaC1}<0.01 \mathrm{MH}_{2} \mathrm{SO}_{4}$
D. $0.01 \mathrm{MH}_{2} \mathrm{~S}<0.01 \mathrm{MNaNO}_{3}<0.01 \mathrm{MNaC1}<0.01 \mathrm{MH}_{2} \mathrm{SO}_{4}$

## Answer: B

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35. pH of water is 7 . When a substance $Y$ is dissolved in water, the $p H$ becomes 13 . The substance $Y$ is a salt of
A. Weak acid and weak base
B. Strong acid and strong base
C. Strong acid and weak base
D. Weak acid and strong base

## Answer: D

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36. The hydrolysis constant of 0.1 M aqueous solution of sodium acetate if $K_{a}$ of $\mathrm{CH}_{3} \mathrm{COOH}=1.8 \times 10^{-5}$ is
A. $5.5 \times 10^{-10}$
B. $4.5 \times 10^{-8}$
C. $5.5 \times 10^{-12}$
D. None of these

## Answer: A

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37. The compound whose 0.1 M solution is basic is
A. $\mathrm{CH}_{3} \mathrm{COONH}_{4}$
B. $\mathrm{NH}_{4} \mathrm{Cl}$
C. $\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4}$
D. $\mathrm{CH}_{3} \mathrm{COONa}$

## Answer: D

## D Watch Video Solution

38. $K_{a}$ for ascorbic acid (HASc) is $5 \times 10^{-5}$. Calculate the $\left[H^{\oplus}\right]$ in an aqueous solution in which the concentration of $A s C$ ( ${ }^{\Theta}$ ions is $0.02 M$.
A. $2 \times 10^{-6}$
B. $2 \times 10^{-7}$
C. $5 \times 10^{-9}$
D. $5 \times 10^{-10}$

## Answer: C

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## Ex 8.4

1. a. At what pH does indicator change colour if the indicator is a weak acid with $K_{\text {Ind }}=4.0 \times 10^{-4}$.
b. For which of the following neutralisation would the indicator be useful?
i. $\mathrm{HCl}+\mathrm{NaOH}$
ii. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{NaOH}$
iii. $\mathrm{HCl}+\mathrm{NH}_{3}$
c. Name the indicators which can be used for such titration.

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2. The acid from of an acid base indicator is yellow in acid and red in basic from. What is the change in $p H$ in order to change the indicator form $80 \%$ yellow to $80 \%$ red.

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3. Given that solubility product of $\mathrm{BaSO}_{4}$ is $1 \times 10^{-10}$ will be precipiate from when

Equal volumes of $2 \times 10^{-3} \mathrm{MBaCl}_{2}$ solution and $2 \times 10^{-4} \mathrm{MNa}_{2} \mathrm{SO}_{4}$ solution, are mixed?

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4. The $K_{s p}$ of AgCl at $25^{\circ} \mathrm{C}$ is $1.6 \times 10^{-9}$, find the solubility of salt in $g L^{-1}$ in water.

## - Watch Video Solution

5. If solutbility of $\mathrm{Ca}\left(\mathrm{IO}_{3}\right)_{2}$ in water at $20^{\circ} \mathrm{C}$ is $3.9 g \mathrm{~L}^{-1}$. Calculate the $K_{s p}$. Given $\mathrm{MwCa}\left(\mathrm{IO}_{3}\right)_{2}=390$.

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6. Find the solubility of $\mathrm{Ca}\left(\mathrm{IO}_{3}\right)_{2}$ is $\mathrm{molL}^{-1}$ in a solution containing 0.1 MCaCl at $25^{\circ} \mathrm{C} . \mathrm{K}_{\text {sp }}$ of $\mathrm{Ca}\left(\mathrm{IO}_{3}\right)_{2}=6.3 \times 10^{-7}$

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7. The solubility product $\left(K_{s p}\right)$ of $\mathrm{BaSO}_{4}$ is $1.5 \times 10^{-9}$. Calculate the solubility of barium sulphate in pure water and in $0.1 \mathrm{M} \mathrm{BaCl}_{2}$.

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8. A solution contains $1.4 \times 10^{-3} \mathrm{MAgNO}_{3}$. What concentration of $\mathrm{KC1}$ will be required to initiate the precipitation of $A g C 1$ ? $K_{s p} A g C 1=2.8 \times 10^{-10}$

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9. If the solubility of $\mathrm{CaSO}_{4}$ in $\mathrm{H}_{2} \mathrm{O}$ is $10^{-5} \mathrm{M}$, Calculate the solubility in 0.005 M solution of $\mathrm{H}_{2} \mathrm{SO}_{4}$.
10. The concentration of $\mathrm{Ag}^{\oplus}$ ions in a saturated solution of $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ is $2.2 \times 10^{-4} \mathrm{M}$. Calculate the solubility product of $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$

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11. 500 mL of $2 \times 10^{-3} \mathrm{MA1C1}_{3}$ and 500 mol of $4 \times 10^{-2} \mathrm{M}$ solution of NaOH are mixed and solution is diluted to $10^{-2} \mathrm{~L}$ with water at room temperature wil a precipiate exist? Given:
$K_{\text {sp }}$ of $A 1(\mathrm{OH})_{3}=5 \times 10^{-33}$.

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12. You are provided with 500 mL of hard water, containing 0.005 mol of $\mathrm{CaC1}_{2}$ and two $\mathrm{H}_{2} \mathrm{SO}_{4}$ samples of 0.001 M and 0.02 M concentration. Which one or both or none can be used for precipitating $\mathrm{Ca}^{2+}$ ions. $K_{s p}$ of $\mathrm{CaSO}_{4}=2.4 \times 10^{-4}$.

## (D) Watch Video Solution

13. A lead salts is dissolved in HC1 which si $94 \%$ ionised. It is found to have $0.1 \mathrm{MPb}^{2+}$ and $0.28 \mathrm{MH}^{\oplus}$ ions. The solution is satured with $\mathrm{H}_{2} \mathrm{~S}(\mathrm{~g})$. Calculate the amount of $\mathrm{Pb}^{2+}$ ions that remains unprecipitated.
$K_{s p}$ of $P b S=4 \times 10^{-29}$,
$K_{\text {sp }}$ of $H_{2} \mathrm{~S}=1.1 \times 10^{-22}$

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14. A solution constains $\mathrm{Zn}^{2+}$ ions and $\mathrm{Cu}^{2+}$ ions each of 0.02 M . If the solution is made $1 M$ in $H^{\oplus}$, and $H_{2} S$ is passed untill the solution is satured, should a precipitate be formed? Given: $K_{s p} Z n S=10^{-22}$,

$$
K_{s p} \text { Cus }=8 \times 10^{-37} .
$$

In satured solution, $K_{s p}\left(H_{2} \mathrm{~S}\right)=10^{-22}$

## - Watch Video Solution

15. The following pH range where the indicator shows change in colour are given
i. 4 - 9.7 ii. $7.46-10.0$ iii. $6.5-4$

Which of the above $p H$ range represent titration of
I. Strong acid/strong base $\left(S_{A} / S_{B}\right)$,
II. Weak acid/strong base $\left(W_{A} / S_{B}\right)$,
III. Weak base/strong acid $\left(W_{B} / S_{A}\right)$
A. (i) $\rightarrow$ I, (ii) $\rightarrow$ II, (iii) $\rightarrow$ III
B. (iii) $\rightarrow$ I, (ii) $\rightarrow$ II, (i) $\rightarrow$ III
C. (ii) $\rightarrow$ I, (iii) $\rightarrow$ II, (i) $\rightarrow$ III
D. (i) $\rightarrow$ I, (iii) $\rightarrow$ II, (ii) $\rightarrow$ III

## Answer: A

16. The following acid base titration graphs are given:

(I)

(II)
(I)


Which of the following graph represents titration of
i. $\mathrm{NH}_{4} \mathrm{OH} / \mathrm{HCl}\left(W_{B} / S_{A}\right)$
ii. $\mathrm{HNO}_{3} / \mathrm{KOH}\left(\mathrm{S}_{\mathrm{A}} / S_{B}\right)$
iii. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOH} / \mathrm{KOH}\left(W_{A} / S_{B}\right)$

| Araph | Titration |
| :--- | :--- |
| A. |  |
| I. II. III | i. ii. iii |
| Graph | Titration |
| B. ${ }_{\text {I. II. III }}$ | ii. iii. $i$ |
| C.Graph | Titration |
| I. II. III | iii. ii. $i$ |

D. $\begin{array}{ll}\text { Graph } & \text { Titration } \\ \text { I. III } & \text { iii. i. ii }\end{array}$

## Answer: B

## D Watch Video Solution

17. What indicators will be suitable for the following acid-base titrations:
a. HCOOH against NaOH
b. HBr against KOH
c. $\mathrm{NH}_{4} \mathrm{OH}$ with $\mathrm{HNO}_{3}$
A. (i) $\rightarrow$ A, (ii) $\rightarrow B$, (iii) $\rightarrow C$
B. (i) $\rightarrow$, (ii) $\rightarrow C$, (iii) $\rightarrow B$
C. (i) $\rightarrow B$, (ii) $\rightarrow C$, (iii) $\rightarrow A$
D. $(\mathrm{i}) \rightarrow C,(\mathrm{iii}) \rightarrow A,($ iii $) \rightarrow B$

## Answer: D

18. A solution gives the following colours with different indicators:
a. Methy1 orange $\Rightarrow$ Yellow
b. Methy1 red $\Rightarrow$ Yellow
c. Bromothymol blue $\Rightarrow$ Orange

What is the pH of the solution?
A. $>4.5$
B. $>6.0$
C. 6.0 to 6.3
D. 4.5 to 6

## Answer: C

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19. In the titration of $\mathrm{NH}_{4} \mathrm{OH}$ versus HCl , the pH of the solution at equivalence point is about:
A. 5.5
B. 7
C. 8.5
D. 9.5

## Answer: A

## D Watch Video Solution

20. The $p H$ indicators are
A. Salts of strong acids and strong bases
B. Salts of weak acids and weak bases
C. Either weak acids or weak bases
D. Either strong acids or strong base

## Answer: C

21. In which of the following acid-base titration, the pH is greater than 8 at the equivalence point?
A. Aceitic acid vs ammonia
B. Acetic acid vs sodium hydroxide
C. Hydrochloric acid vs ammonia
D. Hydrochloric acid vs sodium hydroxide

## Answer: B

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22. Strong acids are generally used as standard solution in acid-base titrations because:
A. The $p H$ at the equivalent point will always be 7 .
B. They can be used to titrate both strong and weak bases.
C. Strong acids from more stable soluitons than weak acids.
D. The salts of strong acids do not hydrolyse.

## Answer: C

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23. The best indicator for detection of end point in titration of a weak acid and a strong base is
A. Methy 1 orange ( 3 to 4 )
B. Methy1 red (5 to 6)
C. Bromotymol blue (6 to 7.5)
D. Phenolphthalein (8 to 9.6)

## Answer: D

24. The precipitate of $\operatorname{CaF}_{2}\left(K_{s p}=1.7 \times 10^{-10}\right)$ is obtained when equal volumes of the following are mixed
A. $10^{-2} \mathrm{MCa}^{2+}+10^{-3} \mathrm{MF}^{\Theta}$
B. $10^{-4} \mathrm{MCa}^{2+}+10^{-4} \mathrm{MF}^{\Theta}$
C. $10^{-3} \mathrm{MCa}^{2+}+10^{-5} \mathrm{MF}^{\Theta}$
D. $10^{-5} \mathrm{MCa}^{2+}+10^{-3} \mathrm{MF}^{\Theta}$

## Answer: A

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25. The solubility of $A_{2} B_{3}$ is "x mol dm"^(-3). ItsK_(sp)' is
A. $6 x^{4}$
B. $64 x^{4}$
C. $36 x^{5}$
D. $108 x^{5}$

## Answer: D

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26. The pH of $\mathrm{Ca}(\mathrm{OH})_{2}$ is 10.6 at $25^{\circ} \mathrm{C} . \mathrm{K}_{s p}$ of $\mathrm{Ca}(\mathrm{OH})_{2}$ is
A. $3.2 \times 10^{-12} M^{3}$
B. $3.2 \times 10^{-11} M^{3}$
C. $1.6 \times 10^{-12} M^{3}$
D. $1.6 \times 10^{-11} M^{3}$

## Answer: B

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27. Solubility of AgI in $0.05 \mathrm{MBaI}_{2}$ solution is $10^{-15} \mathrm{M}$. The solubility of AgI in water is
A. $25 \times 10^{-7}$
B. $10^{-7} \mathrm{M}$
C. $5 \times 10^{-8}$
D. $10^{-8} \mathrm{M}$.

## Answer: D

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28. Solubility of a solute in water is dependent on temperature as given by
$S=A e^{-\Delta H / R T}$, where $\Delta H=$ heat of solution
Solute $+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow$ Solution, $\Delta H= \pm x$
For given solution, variation of $\log \mathrm{S}$ with temperature is shown
graphically. Hence, solution is

A. CaO
B. $\mathrm{MgSO}_{4}$
C. $\mathrm{CuSO}_{4}$
D. $\mathrm{CuSO}_{4} \cdot 5 \mathrm{H}_{2} \mathrm{O}$
29. The solubility of $\mathrm{CaF}_{2}$ in a solution of $0.1 \mathrm{MCa}\left(\mathrm{NO}_{3}\right)_{2}$ is
A. $\left[\mathrm{Ca}^{2+}\right]$
B. $2\left[F^{\Theta}\right]$
$\left[F^{\Theta}\right]$
C. $\frac{}{2}$
D. $2\left[\mathrm{NO}_{3}^{\Theta}\right]$

## Answer: C

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30. The volume of water needed to dissolve 1 mg of $\mathrm{PbSO}_{4}\left(K_{\text {sp }}=1.44 \times 10^{-8}, \mathrm{MwofPbSO}_{4}=303 \mathrm{~g}\right)$ at $25^{\circ} \mathrm{C}$ is
A. 80 mL
B. $43 m L$
C. 27.5 mL
D. 10 mL

## Answer: C

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31. The volume of water needed to prepare a satured solution of $\mathrm{Ag}^{\oplus}$ having maximum $\left[A g^{\oplus}\right]$ ion by selecting one out of three slats form:
$\operatorname{AgC1}\left(K_{s p}=2.0 \times 10^{-10}\right), \operatorname{AgBr}\left(K_{s p}=5 \times 10^{-13}\right)$,
and
$\mathrm{Ag}_{2} \mathrm{CrO}_{4}\left(\mathrm{~K}_{\text {sp }}=2.4 \times 10^{-12}\right)$. whcih compound should be used to have maximum $\left[A g^{\oplus}\right]$ ?
A. $\mathrm{AgC1}$
B. AgBr
C. $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$
D. Any one of them

## Answer: C

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32. How many grams of KBr can be added to 1 L of 0.12 M solution of $\mathrm{AgNO}_{3}$ just to start the precipitation of
$\operatorname{AgBr} .\left(M w o f K B r=120, K_{s p} o f A g B r=10^{-13}\right)$
A. $10^{-10} g$
B. $10^{-9} \mathrm{~g}$
C. $0.5 \times 10^{-10} g$
D. $0.5 \times 10^{-9} g$

## Answer: A

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33. The solubility of silver benzoate $\left(\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOAg}\right)$ in $\mathrm{H}_{2} \mathrm{O}$ and in a buffer solution of $p H=2,3$, and 4 are $S_{1}, S_{2}, S_{3}$ and $S_{4}$ respectively. The decreasing order of solubility is
A. $S_{1}>S_{2}>S_{3}>S_{4}$
B. $S_{4}>S_{3}>S_{2}>S_{1}$
C. $S_{2}>S_{3}>S_{4}>S_{1}$
D. $S_{3}>S_{2}>S_{4}>S_{1}$

## Answer: C

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34. The solubility of $\mathrm{CH}_{3} \mathrm{COOAg}$ in a buffer solution with $\mathrm{pH}=4$, whose $K_{s p}=10^{-12}$ and $K_{a}=\frac{10^{-4}}{3}$ is
A. $10^{-6}$
B. $0.5 \times 10^{-6}$
C. $5 \times 10^{-6}$
D. $2 \times 10^{-6}$

## Answer: D

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35. the ratio of solubility of $\mathrm{CH}_{3} \mathrm{COOAg}$ in a buffer solution with $\mathrm{pH}=4$ and in $\mathrm{H}_{2} \mathrm{O}$ is
(SolBuffer $\left.=2 \times 10^{-6}\right)$
A. $1 / 2$
B. 2
C. $1 / 3$
D. 3

## Answer: B

36. What is the maximum molarity of $\mathrm{Co}^{+2}$ ions in $0.1 \mathrm{MHC1}$ saturated with $0.1 \mathrm{MH}_{2} S .\left(K_{a}=4 \times 10^{-21}\right)$. Given: $K_{s p}$ of $\operatorname{CoS}=2 \times 10^{-21}$.
A. 0.10 M
B. 1.00 M
C. $4.48 \times 10^{-11} M$
D. 0.050 M

## Answer: D

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37. The following curve shows the change of pH during the course of titration of weak acid HA with a strong base. At which point in the titration curve is the concentration of acid equal to that of its conjugate
base.

A. Point $B$
B. Point $C$
C. Point $D$
D. Point $E$

Answer: C
38. If the salts $M_{2} X, Q Y_{2}$, and $P Z_{3}$ have the same solubilities $\left(<\frac{4}{27}\right)$, their $K_{s p}$ values are related
A. $K_{s p}\left(M_{2} X\right)=K_{s p}\left(Q Y_{2}\right)>K_{s p}\left(P Z_{3}\right)$
B. $K_{s p}\left(M_{2} X\right)>K_{s p}\left(Q Y_{2}\right)=K_{s p}\left(P Z_{3}\right)$
C. $K_{s p}\left(M_{2} X\right)=K_{s p}\left(Q Y_{2}\right)=K_{s p}\left(P Z_{3}\right)$
D. $K_{s p}\left(M_{2} X\right)>K_{s p}\left(Q Y_{2}\right)>K_{s p}\left(P Z_{3}\right)$

## Answer: A

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39. Arrange the following solutions in decreasing order of $\left[\mathrm{Ag}^{\oplus}\right]$ ion:
I. $1 M\left[\operatorname{Ag}(C N)_{2}\right]^{\Theta}$
II. Saturated $\mathrm{AgC1}$
III. $1 M\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{\oplus} \in 0.1 \mathrm{MNH}_{3}$
IV. Saturated AgI

$$
\begin{aligned}
& \left(K_{s p} o f A g C 1=10^{-10}, K_{s p} o f A g I=8.3 \times 10^{-17} K_{f}\right. \\
& {\left[A g\left(C N_{2}\right)\right]^{\Theta}=10^{21}, K_{f}\left[A g\left(\mathrm{NH}_{3}\right)_{2}\right]^{\oplus}=10^{8}}
\end{aligned}
$$

A. $I>I I>I I I>I V$
B. II $>$ III $>I>I V$
C. IV $>$ II $>$ II $>I$
D. $I>I V>I I I>I I$

## Answer: B

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## Ex 8.5

1. How many moles of $\mathrm{NH}_{3}$ must be added to 1.0 L of $0.75 \mathrm{MAgNO}_{3}$ in order to reduce the $\left[\mathrm{Ag}^{\oplus}\right]$ to $5.0 \times 10^{-8} \mathrm{M} . \mathrm{K} \mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{\oplus}=1 \times 10^{8}$.
2. Calculate the $\left[\mathrm{Fe}^{2}\right]$ in a solution prepared by mixting 75.0 mL of $0.03 \mathrm{MFeSO}_{4}$ with 125.0 mL of $0.2 \mathrm{M} \mathrm{KCNK} \mathrm{Fe}_{\mathrm{Fe}}(\mathrm{CN})_{6}^{4-}=1 \times 10^{24}$.

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3. a. Calculate $\left[\mathrm{Ag}^{\oplus}\right]$ in a solution of $\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{\oplus}\right]$ prepared by adding
$1.0 \times 10^{-3} \mathrm{molAgNO}_{3}$ to $1.0 \mathrm{Lof1.0MNH}_{3}$ solution $\mathrm{Kf} \mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{\oplus}=10^{8}$.
b. Calculate $\left[\mathrm{Ag}^{\oplus}\right]$ which is in equilibrium with $0.15 \mathrm{M}\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{\oplus}$ and $1.5 \mathrm{NH}_{3}$.

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4. Calculate the $\left[\mathrm{Fe}^{2+}\right]$ in a solution containing $0.2 \mathrm{M}\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{4-}$ and $0.1 M C N^{\Theta} \cdot K_{f} \mathrm{Fe}(C N)_{6}^{4-}=1 \times 10^{24}$.

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5. How much AgBr could dissolve in 1.0 L of $0.4 \mathrm{MNH}_{3}$ ? Assume that $\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{\oplus}$ is the only complex formed. Given: the dissociation constant for
$\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{\oplus} \Leftrightarrow \mathrm{Ag}^{\oplus}+2 \mathrm{NH}_{3}$,
$K_{d}=6.0 \times 10^{-8}$ and $K_{\text {sp }}(\mathrm{AgBr})=5.0 \times 10^{-13}$.

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6. Calculate $K_{f}$ for the reaction:
$M^{3+}+S C N^{\Theta} \Leftrightarrow M S C N^{2+}$,
The $\left[M^{3+}\right]$ in the solution is $2.0 \times 10^{-3} \mathrm{M},\left[S C N^{\Theta}\right]=1.5 \times 10^{-3} \mathrm{M}$ and
Free $\left[S C N^{\Theta}\right]=1.0 \times 10^{-5} \mathrm{M}$.

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1. a. Distinguish between acid strength and acid concentration.
b. Distinuish between weak base and an isoluble base.

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2. Write an equilibrium equation for a solution containing $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COONa}$. What effect does $\mathrm{CH}_{3} \mathrm{COONa}$ have on a solution of $\mathrm{CH}_{3} \mathrm{COOH}$ ?

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3. Which of the reagents listed below could be added to water to make
0.1 M solutions of each of the following ions?
$\oplus$
a. $\mathrm{NH}_{4}$
b. $\mathrm{CH}_{3} \mathrm{COO}^{\Theta}$
c. $C 1^{\Theta}$
i. $\mathrm{NH}_{3}$ ii. $\mathrm{NH}_{4} \mathrm{Cl}$
iii. $\mathrm{CH}_{3} \mathrm{COOH}$
iv. $\mathrm{CH}_{3} \mathrm{COONa}$
v. HC 1
vi. $N a C 1$.

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4. Saccharin $\left(K_{a}=2 \times 10^{-12}\right)$ is a weak acid represented by formula HSaC. A $4 \times 10^{-4}$ mole amount of saccharin is dissolved in $200 \mathrm{~cm}^{3}$ water of pH 3 . Assuming no change in volume. Calculate the soncentration of $S a C^{-}$ions in the resulting solution at equilibrium.

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5. Accetyl salicylic acid (aspirin) ionises in water as:


$$
\left(K_{a}=2.75 \times 10^{-9}\right)
$$

If two tablets of aspirin each of 0.32 g is dissolved in water to produce 250 mL solution, calculate


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6. Calculate the $\left[\mathrm{CH}_{2} \mathrm{FCOOH}\right]$ (fluoroacetic acid) which is required to get $\left[H^{\oplus}\right]=1.5 \times 10^{-3} \mathrm{M} . K_{a}$ of acid $=2.6 \times 10^{-3}$.

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7. Calculate the dissociation constant of $\mathrm{NH}_{4} \mathrm{OH}$ at 298 k , if $\Delta H^{\Theta}$ and $\Delta S^{\Theta}$ for the given changes are as follows:-
$N H_{3}+H^{\oplus} \stackrel{\oplus}{\Leftrightarrow} \mathrm{NH}_{4}$,

$$
\Delta H^{\Theta}=-52.2 \mathrm{KJmol}^{-1}, \Delta S^{\Theta}=1.67 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}
$$

$\Theta$
$\mathrm{H}_{2} \mathrm{O} \Leftrightarrow H^{\oplus}+\mathrm{OH}, \Delta H^{\Theta}=56.6 \mathrm{kJmol}^{-1}$.
$\Delta S^{\Theta}=-76.53 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}$

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8. Determine the dergee of dissociation of $0.05 \mathrm{MNH}_{3}$ at $25^{\circ} \mathrm{C}$ in a solution of $\mathrm{pH}=11$

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9. In the eqantitative analysis $\mathrm{Bi}^{3+}$ is detected precipitation of $[\mathrm{BiO}(\mathrm{OH})(\mathrm{s})]$ [bismuthy1 hydroxide). Calculate the pH when the following
equilibria exists:
$\mathrm{BiO}(\mathrm{OH})(\mathrm{s}) \Leftrightarrow \mathrm{BiO}^{\oplus}(a q)+\stackrel{\Theta}{\mathrm{O}} \mathrm{H}(\mathrm{aq})\left(\mathrm{K}=4 \times 10^{-10}\right)$

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10. Calculate $\left[\begin{array}{l}\Theta \\ \mathrm{OH}\end{array}\right]$ of $\left[\mathrm{NH}_{2} \mathrm{C}_{2} \mathrm{H}_{4} \mathrm{NH}_{3}\right]^{\oplus}$ and $\left[\mathrm{H}_{3} \mathrm{~N}-\mathrm{C}_{2} \mathrm{H}_{4} \mathrm{NH}_{3}\right]^{2+}$ in
$0.15 M$ ethylene diamine (aq) if
$\mathrm{NH}_{2} \mathrm{C}_{2} \mathrm{H}_{4} \mathrm{NH}_{2}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{NH}_{2} \mathrm{C}_{2} \mathrm{H}_{4} \stackrel{\oplus}{\mathrm{~N}} \mathrm{H}_{3}+\stackrel{\ominus}{\mathrm{O}} \mathrm{H}\left(\mathrm{K}_{1}=8.5 \times 10^{-5}\right)$
$\mathrm{NH}_{2} \mathrm{C}_{2} \mathrm{H}_{4} \stackrel{\oplus}{\mathrm{~N}} \mathrm{H}_{3}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow\left[\mathrm{NH}_{3} \mathrm{C}_{2} \mathrm{H}_{4} \mathrm{NH}_{3}\right]^{2+}+\stackrel{\ominus}{\mathrm{O}} \mathrm{H}\left(\mathrm{K}_{2}=2.7 \times 10^{-8}\right)$

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11. Calculate pH of
a. $0.002 \mathrm{NCH}_{3} \mathrm{COOH}$ having $2.3 \%$ dissociation.
b. $0.002 \mathrm{NNH}_{4} \mathrm{OH}$ having $2.3 \%$ dissociation.

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12. Calculate $\left[\mathrm{H}^{\oplus}\right]$ and $\left[\mathrm{CHCl}_{2} \mathrm{COO}^{\Theta}\right]$ in a solution that is 0.01 MHCl and $0.01 \mathrm{MCHCl}_{2} \mathrm{COOH}$. $\mathrm{K}_{a}$ for $\mathrm{CHCl}_{2} \mathrm{COOH}$ is $5 \times 10^{-3}$.

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13. A solution contains $0.09 \mathrm{HC}, 0.09 \mathrm{MCHC1}_{2} \mathrm{COOH}$, and $0.1 \mathrm{MCH}_{3} \mathrm{COOH}$. The pH of this solution is one. Calculate $\mathrm{K}_{a}$ for $\mathrm{CHC1}_{2} \mathrm{COOH}$. (Given $\left.\mathrm{K}_{a} \mathrm{CH}_{3} \mathrm{COOH}=10^{-5}\right)$

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14. What is the concentration of $\mathrm{CH}_{3} \mathrm{COOH}$ which can be added to 0.5 MHCOOH solution so that dissociation of both is same.

$$
K_{\mathrm{CH}_{3} \mathrm{COOH}}=1.8 \times 10^{-5}, K_{\mathrm{HCOOH}}=2.4 \times 10^{-4}
$$

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15. What are $\left[H^{\oplus}\right],\left[A^{\Theta}\right.$, and $\left[B^{\Theta}\right]$ in a solution that is $0.3 M H A$ and $0.1 \mathrm{MHB} ? K_{a}$ for $H A$ and $H B$ are $1.38 \times 10^{-4}$ and $1.05 \times 10^{-10}$, respectively.

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## Exercises Subjective (Buffer Solutions)

1. What will be the amount of $\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4}$ (in g) which must be added to 500 mL of $0.2 \mathrm{M} \mathrm{NH} \mathrm{H}_{4} \mathrm{OH}$ to yield a solution of pH 9.35 ?
$\left[\right.$ Given, $p K_{a}$ of $\left.\mathrm{NH}_{4}^{+}=9.26, p K_{b} \mathrm{NH}_{4} \mathrm{OH}=14-p K_{a}\left(\mathrm{NH}_{4}^{+}\right)\right]$

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2. A 0.1 Msolution of weak acid $H A$ is $1 \%$ dissociated at $298 k$. what is its $K_{a}$ ? what will be the new degree of dissociation of $H A$ and $p H$ when $0.2 M$ of $N a A$ is added to it.

## Exercises Subjective (Hydrolysis Of Salt)

1. a. Calculate the percentage hydrolysis of 0.003 M aqueous solution of $\mathrm{NaOH} . \mathrm{K}_{a}$ for $\mathrm{HOCN}=3.3 \times 10^{-4}$.
b. What is the $p H$ and $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$ of $0.02 M$ aqueous solution of sodium butyrate. $\left(K_{a}=2.0 \times 10^{-5}\right)$.

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2. $K_{a}$ for the reaction, $\mathrm{Fe}^{3+}($ aq. $)+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{Fe}(\mathrm{OH})^{2+}($ aq. $)+\mathrm{H}_{3} \mathrm{O}^{+}($aq. $)$ is $6.5 \times 10^{-3}$ What is the max. pH value which could be used so that at least $80 \%$ of the total iron (III) in a dilute solution exists as $\mathrm{Fe}^{3+}$ ?

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1. Calculate the equilibrium constants for the reactions with water of $\mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}, \mathrm{HPO}_{4}^{2-}$, and $\mathrm{PO}_{4}^{3-}$ as ase. Comparing the relative values of two equilibrium constants of $\mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}$ with water, deduce whether solutions of this ion in water are acidic or base, Deduce whether solutions of $\mathrm{HPO}_{4}^{2-}$ are acidic or bases. Given $K_{1}, K_{2}$, and $K_{3}$ for $H_{3} P O_{4}$ are $7.1 \times 10^{-3}, 6.3 \times 10^{-8}$, and $4.5 \times 10^{-13}$ respectively.

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2. Citric acid $\left(\mathrm{H}_{3} A\right)$ is a polyprotic acid with $K_{1}, K_{2}$, and $K_{3}$ equals to $7.4 \times 10^{-4}, 1.7 \times 10^{-5}$, and $4.0 \times 10^{-7}$, respectively. Calculate the $\left[H^{\oplus}\right],\left[H_{2} A^{\Theta}\right],\left[H A^{2-}\right]$, and $\left[A^{3-}\right]$ in $0.01 M$ citric acid.

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## Exercises Subjective (Solubility And $K_{S p}$ )

1. a. 25 mL of sample of saturated solution of $\mathrm{PbI}_{2}$ requires 10 mL of a certain $\mathrm{AgNO}_{3}(a q)$ for its titration. What is the molarity of this $\mathrm{AgNO}_{3}(a q) ? \mathrm{~K}_{\text {sp }}$ of $\mathrm{PbI}_{2}=4 \times 10^{-9}$.
b. $M(\mathrm{OH}) \times$ has $K_{s p}=27 \times 10^{-12}$ and solubility in water is $10^{-3} \mathrm{M}$. Calculate the value of $x$.

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2. Equal volumes of $0.02 \mathrm{MCaC1}_{2}$ and $0.0004 \mathrm{MNa}_{2} \mathrm{SO}_{4}$ are mixed. Will a precipitate form? $K_{\text {sp }}$ for $\mathrm{CaSO}_{4}=2.4 \times 10^{-5}$ ?

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3. What $\left(\mathrm{H}_{3} \mathrm{O}^{+}\right)$must be maintained in a saturated $\mathrm{H}_{2} \mathrm{~S}$ solution to precipitate $\mathrm{Pb}^{2+}$, but not $\mathrm{Zn}^{2+}$ from a solution in which each ion is present at a concetration of 0.01 M ? $\left(K_{S P}\right.$ for $H_{2} S=1.1 \times 10^{-22}, K_{S P}$ for $\mathrm{ZnS}=1.0 \times 10^{-21}$ )
4. Calculate the solubility of $\mathrm{CaF}_{2}$ in a solution buffered at $\mathrm{pH}=3.0 . \mathrm{K}_{a}$ for $H F=6.3 \times 10^{-4}$ and $K_{s p}$ of $C a F_{2}=3.45 \times 10^{-11}$.

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5. Will a precipitate of $\mathrm{Mg}(\mathrm{OH})_{2}$ be formed in a 0.002 M solution of $\mathrm{Mg}\left(\mathrm{NO}_{3}\right)_{2}$ if the pH of solution is adjusted to $9 . K_{\text {sp }}$ of $\mathrm{Mg}(\mathrm{OH})_{2}=8.9 \times 10^{-12}$.

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6. 0.01 mole of $\mathrm{AgNO}_{3}$ is added to 1 litre of a solution which is 0.1 M in $\mathrm{Na}_{2} \mathrm{CrO}_{4}$ and 0.005 M in $\mathrm{NaIO}_{3}$. Calculate the mole of precipitate formed at equilibrium and the concentrations of $\mathrm{Ag}^{+}, \mathrm{IO}_{3}^{-}$and $\mathrm{CrO}_{4}^{2-} .\left(\mathrm{K}_{\mathrm{SP}}\right.$ values of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ and $\mathrm{AgIO}_{3}$ are $10^{-8}$ and $10^{-13}$ respectively)
7. 1.75 g of solid NaOH is added to $0.25 \mathrm{dm}^{3}$ of $0.1 \mathrm{MNiCl}_{2}$ solution.

Calculate:
a. Mass of $\mathrm{Ni}(\mathrm{OH})_{2}$ forms
b. $p H$ if final solution Given $K_{s p}$ of $\mathrm{Ni}(\mathrm{OH})_{2}=1.6 \times 10^{-14}$

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8. Zn salt is mixed with $\left(\mathrm{NH}_{4}\right)_{2} \mathrm{~S}$ of 0.021 M . What amount of $\mathrm{Zn}^{2+}$ will remain uprecipitated in 12 mL of the solution? $K_{s p}$ of $Z n S=4.51 \times 10^{-24}$.

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9. A solution has $0.05 \mathrm{MMg}^{2+}$ and $0.05 \mathrm{MNH}_{3}$. Calculate the concentration of $\mathrm{NH}_{4} \mathrm{CI}$ required to prevent the formation of $\mathrm{Mg}(\mathrm{Oh})_{2}$ in solution. $\mathrm{K}_{S P}$ for $\mathrm{Mg}(\mathrm{OH})_{2}=9.0 \times 10^{-12}$ and ionisation constant of $\mathrm{NH}_{3}$ is $1.8 \times 10^{-5}$.
10. A hard water sample has $131 \mathrm{ppm} \mathrm{CaSO}_{4}$. What fraction of the water must be evaporated in a container before solid $\mathrm{CaSO}_{4}$ begins to deposit. $K_{s p} o f$ CaSO $_{4}=9.0 \times 10^{-6}$.

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11. To a solution of $0.01 \mathrm{MMg}^{2+}$ and $0.8 \mathrm{MNH}_{4} \mathrm{CI}$, and equal volume of $\mathrm{NH}_{3}$ is added which just gives precipitates. Calculate $\left[\mathrm{NH}_{3}\right]$ in solution.

$$
K_{s p} o f M g(\mathrm{OH})_{2}=1.4 \times 10^{-11} \text { and } K_{b} o f \mathrm{NH}_{4} \mathrm{OH}=1.8 \times 10^{-5} .
$$

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12. 10 mL of $0.3 \mathrm{MNa}_{2} \mathrm{SO}_{4}$ are mixed with 20 mL solution having initially $0.1 \mathrm{MCa}^{2+}$ and $0.1 \mathrm{MSr}^{2+}$ in it. Calculate the final $\left[\mathrm{Ca}^{2+}\right],\left[\mathrm{Sr}^{2+}\right]$ and $\left[\mathrm{SO}_{4}^{2-}\right]$ in solution? Given $K_{s p} \mathrm{SrSO}_{4}=7.6 \times 10^{-7}$ and $K_{\text {sp }} \mathrm{CaSO}_{4}=2.4 \times 10^{-5}$.
13. The solubility of $\mathrm{CaCO}_{3}$ is $7 \mathrm{mg} / \mathrm{L}$. Calculate the $\mathrm{K}_{\text {sp }}$ of $\mathrm{BaCO}_{3}$ when $\mathrm{Na}_{2} \mathrm{CO}_{3}$ is added slowly a solution containing equimolar concentration of $\mathrm{Ca}^{2+}$ and $\mathrm{Ba}^{2+}$ and no precipitate is formed until $90 \%$ of $\mathrm{Ba}^{2+}$ has been precipitated as $\mathrm{BaCO}_{3}$.

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14. Calculate the solubility of AgCN in a buffer solution of $p H 3.0$. Assume that no cyano complex is formed $K_{s p} A g C N=2.2 \times 10^{-16}, K_{a} H C N=6.2 \times 10^{-10}$.

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15. Equal volumes of $0.02 \mathrm{MAgNO}_{3}$ and 0.01 MHCN are mixed. Calculate $\left[A g^{\oplus}\right]$ in solution after attaining equilibrium. $K_{a} H C N=6.2 \times 10^{-10}$ and $K_{s p}$ of $A g C N=2.2 \times 10^{-16}$.
16. Determine the number of mole of AgI which may be dissolved in 1.0 litre of $1 M C N^{-}$solution. $K_{S P}$ for AgI and $K_{C}$ for $\mathrm{Ag}(\mathrm{CN})_{2}^{-}$are $1.2 \times 10^{-17} M^{2}$ and $7.1 \times 10^{19} M^{-2}$ respectively.

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17. 100.0 mL of a saturated solution of $\mathrm{Ag}_{2} \mathrm{SO}_{4}$ is added to 250.0 mL of saturated solution of $\mathrm{PbCrO}_{4}$. Will may precipitate form and if so what? Given $\quad K_{s p} \quad$ for $\mathrm{Ag}_{2} \mathrm{SO}_{4}, \mathrm{Ag}_{2} \mathrm{CrO}_{4}, \mathrm{PbCrO}_{4}$, and $\mathrm{PbSO}_{4}$ are $1.4 \times 10^{-5}, 2.4 \times 10^{-12}, 2.8 \times 10^{-13}$, and $1.6 \times 10^{-8}$, respectively.

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18. 2 M solution of $\mathrm{Na}_{2} \mathrm{CO}_{3}$ is boiled in a closed container with excess of $\mathrm{CaF}_{2}$. Very small amount of $\mathrm{CaCO}_{3}$ and NaF are formed. If $\mathrm{K}_{\text {sp }}$ of $\mathrm{CaCO}_{3}$ is $x$ and molar solubility of $\mathrm{CaF}_{2}$ is $y$, find the molar after cocentration of $F^{\Theta}$ in the resulting solution after equilibrium is attained.

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19. $\mathrm{K}_{\text {sp }}$ of $\mathrm{SrF}_{2}=2.8 \times 10^{-9}$ at $25^{\circ} \mathrm{C}$. How much NaF should be added to 100 mL of solution having 0.016 M in $\mathrm{Sr}^{2+}$ ions to reduce its concentration to $2.5 \times 10^{-3} \mathrm{M}$ ?

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## Exercises Subjective(Coordination Equilibria)

1. $\mathrm{H}_{2} \mathrm{~S}$ is bubbled into a 0.2 MNaCN solution which is 0.02 M each in $\mathrm{Ag}(\mathrm{CN})_{2}^{\Theta}$ and $\mathrm{Cd}(\mathrm{CN})_{4}^{2-}$. If $K_{s p}$ of $\mathrm{Ag}_{2} \mathrm{~S}$ and $C d S$ are $10^{-50}$ and $7.1 \times 10^{-28}$ and $K$ instability for $\left[\operatorname{Ag}(C N)_{2}^{\Theta}\right]$ and $\left[C d(C N)^{2-1} 4\right]$ are $1.0 \times 10^{-20}$ and $7.8 \times 10^{-18}$, which sulphide will precipitate first?

## - Watch Video Solution

2. Calculate the equilibrium constants of each of the indicated species necessary to reduce an initial $0.2 \mathrm{MZn}^{2+}$ solution to $1.0 \times 10^{-4} \mathrm{Zn}^{2+}$.
$\Theta$
OH in equilibrium with $\mathrm{Zn}(\mathrm{OH})_{2}(\mathrm{~s})$.
Given.
$K_{s p} Z N(O H)_{2}=1.8 \times 10^{-14}$.
$K_{f} \mathrm{Zn}(\mathrm{OH})_{4}^{2-}=5 \times 10^{14}$.

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## Exercises Linked Comprehension

1. $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$ is a sparingly soluble salt $\left(K_{s p}=2.6 \times 10^{-13}\right)$. To 35 mL of $0.15 \mathrm{MPb}\left(\mathrm{NO}_{3}\right)_{2}$ solution, 15 mL of $0.8 \mathrm{MKIO}_{3}$ solution is added, and a precipiatte of $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$ is formed.

Which is the limiting reactant of teh reaction that takes place in the solution?
A. $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$
B. $\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}$
C. $\mathrm{KIO}_{3}$
D. Both (b) and (c).

## Answer: B

## - Watch Video Solution

2. $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$ is a sparingly soluble salt $\left(K_{\text {sp }}=2.6 \times 10^{-13}\right)$. To 35 mL of $0.15 \mathrm{MPb}\left(\mathrm{NO}_{3}\right)_{2}$ solution, 15 mL of $0.8 \mathrm{MKIO}_{3}$ solution is added, and a precipiatte of $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$ is formed.

What will be molarity of $\mathrm{Pb}^{2+}$ ions in the solution after completion of the reactions? $\left[\mathrm{IO3}^{-}\right]=0.03 \mathrm{M}$
A. 0.152
B. 0.081
C. 0.41
D. 0.03

## Answer: D

## D Watch Video Solution

3. $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$ is a sparingly soluble salt $\left(K_{\text {sp }}=2.6 \times 10^{-13}\right)$. To 35 mL of $0.15 \mathrm{MPb}\left(\mathrm{NO}_{3}\right)_{2}$ solution, 15 mL of $0.8 \mathrm{MKIO}_{3}$ solution is added, and a precipiatte of $\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}$ is formed.

What will be molarity of $\mathrm{Pb}^{2+}$ ions in the solution after completion of the reactions? $\left[\mathrm{IO}^{-}\right]=0.03 \mathrm{M}$
A. $8.4 \times 10^{-10}$
B. $1.6 \times 10^{-10}$
C. $2.8 \times 10^{-10}$
D. $6.1 \times 10^{-10}$

## Answer: C

4. Acid-base indicator such as methy1 orange, phenolphthalein, and bromothymol blue ate substances which change colour accroding to the hydrogen ion concentration of the solution to which they are added.

Most indicators are weak acids (or more rarely weak base) in which the undissociated and dissociated forms have different and distinct colours.

If methy 1 orange is used as the examples and the un-dissociated forms is written as $H M O$, then dissociation occurs as shown below:
$H M O \Leftrightarrow H^{\oplus}+\quad M O^{\ominus}$
Reaction:
Red Colourless Yellow
The indicator should have a sharp colour change with the equivalence point of the titration. Usually the colour change of the indicator occurs over a range of about two pH units. It should be noted that the eye cannot detect the exact end point of the tiytration. The $p K_{a}$ of the indicator should be near the pH of the solution at the equivalance point.

Which of the following sitution exists at the equivalence point of titration?
A. $\left[H^{\oplus}\right]=10^{-7} M$
B. $\left[H^{\oplus}\right]=\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$
C. $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]=10^{-7} M$
D. $\frac{\left[H^{\oplus}\right]}{}=10^{-14}$
$\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$

## Answer: B

## - Watch Video Solution

5. If methyl orange is used as a solution at $\mathrm{pH}=2$ containing the indicator would be
A. Orange
B. Yellow
C. Colorless
D. Red

Answer: D
6. Acidic solution is defined as a solution whose $\left[H^{\oplus}\right]>\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$. Base solution has $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]>\left[H^{\oplus}\right]$. During acid-base titrations, $p H$ of the mixture will change depending on the amount base added. This variation is shown in the form of graph by making plot as titration curves 100 mL of $1.0 \mathrm{MH}_{3} \mathrm{~A}\left(K_{a_{1}}=10^{-3}, K_{a_{2}}=10^{-5}, K_{a_{3}}=10^{-7}\right)$ is titrated against 0.1 MNaOh . The titration curve is as follows.


What is the $p H$ at point ${ }^{\wedge}$ ?
A. 3
B. 4
C. 5
D. 6

## Answer: B

## - Watch Video Solution

7. Acidic solution is defined as a solution whose $\left[H^{\oplus}\right]>\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$. Base solution has $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]>\left[H^{\oplus}\right]$. During acid-base titrations, $p H$ of the mixture will change depending on the amount base added. This variation is shown in the form of graph by making plot as titration curves 100 mL of $1.0 \mathrm{MH}_{3} \mathrm{~A}\left(K_{a_{1}}=10^{-3}, K_{a_{2}}=10^{-5}, K_{a_{3}}=10^{-7}\right)$ is titrated against 0.1 MNaOh . The titration curve is as follows.


What is the pH at point ${ }^{\wedge} \mathrm{A}$ ?
A. 11.0
B. 10.2
C. 9.7
D. 7.7

Answer: C
8. Acidic solution is defined as a solution whose $\left[H^{\oplus}\right]>\left[\begin{array}{l}\Theta \\ O H\end{array}\right]$. Base solution has $\left[\begin{array}{l}\Theta \\ O H\end{array}\right]>\left[H^{\oplus}\right]$. During acid-base titrations, $p H$ of the mixture will change depending on the amount base added. This variation is shown in the form of graph by making plot as titration curves 100 mL of $1.0 \mathrm{MH}_{3} A\left(K_{a_{1}}=10^{-3}, K_{a_{2}}=10^{-5}, K_{a_{3}}=10^{-7}\right)$ is titrated against 0.1 MNaOh . The titration curve is as follows.


What will be the change in $p H$ from point $B$ to point $C$ ?
A. 2.8
B. 3.2
C. 4.6

## Answer: D

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9. In equalitative analysis, cations of graph II as well as group IV both are precipitated in the form of sulphides. Due to low value of $K_{s p}$ of group II sulphides, group reagent is $\mathrm{H}_{2} \mathrm{~S}$ in the presence of dil. $\mathrm{HC1}$, and due to high value of $K_{s p}$ of group $I V$ sulphides, group reagent is $H_{2} S$ in the presence of $\mathrm{NH}_{4} \mathrm{OH}$ and $\mathrm{NH}_{4} \mathrm{C}$. In a solution containing 0.1 M each of $\mathrm{Sn}^{2+}, \mathrm{Cd}^{2+}$, and $\mathrm{Ni}^{2+}$ ions, $\mathrm{H}_{2}$ Sgas is passed.
$K_{s p} o f S n S=8 \times 10^{-29}, K_{s p} o f C d S=1510^{-28}, K_{s p} o f N i S-3 \times 10^{-21}, K_{1} o f H_{2} S=1 \times$ If $\mathrm{H}_{2} \mathrm{~S}$ is passed into the above mixture in the presence of HC 1 , which ion will be precipitated first?
A. SnS
B. $C d S$
C. NiS
D. SnS and CdS (both together)

## Answer: C

## - Watch Video Solution

10. In equalitative analysis, cations of graph II as well as group IV both are precipitated in the form of sulphides. Due to low value of $K_{s p}$ of group II sulphides, group reagent is $\mathrm{H}_{2} \mathrm{~S}$ in the presence of dil. $\mathrm{HC1}$, and due to high value of $K_{s p}$ of group $I V$ sulphides, group reagent is $\mathrm{H}_{2} \mathrm{~S}$ in the presence of $\mathrm{NH}_{4} \mathrm{OH}$ and $\mathrm{NH}_{4} \mathrm{C} 1$. In a solution containing 0.1 M each of $\mathrm{Sn}^{2+}, \mathrm{Cd}^{2+}$, and $\mathrm{Ni}^{2+}$ ions, $\mathrm{H}_{2}$ Sgas is passed.
$K_{s p} o f S n S=8 \times 10^{-29}, K_{s p}$ ofCdS $=1510^{-28}, K_{s p} o f N i S-3 \times 10^{-21}, K_{1} o f H_{2} S=1 \times$
At what value of pH , NiS will start to precipitate?
A. 12.76
B. 7
C. 1.24
D. $4^{\prime}$

## D Watch Video Solution

11. In equalitative analysis, cations of graph $I I$ as well as group $I V$ both are precipitated in the form of sulphides. Due to low value of $K_{s p}$ of group II sulphides, group reagent is $H_{2} S$ in the presence of dil. $H C 1$, and due to high value of $K_{s p}$ of group $I V$ sulphides, group reagent is $H_{2} S$ in the presence of $\mathrm{NH}_{4} \mathrm{OH}$ and $\mathrm{NH}_{4} \mathrm{C} 1$. In a solution containing 0.1 M each of $\mathrm{Sn}^{2+}, \mathrm{Cd}^{2+}$, and $\mathrm{Ni}^{2+}$ ions, $\mathrm{H}_{2}$ Sgas is passed. $K_{s p} o f S n S=8 \times 10^{-29}, K_{s p} o f C d S=1510^{-28}, K_{s p} o f N i S-3 \times 10^{-21}, K_{1} o f H_{2} S=1 \times$

Which of the following sulphides is more soluble in pure water?
A. $C d S$
B. NiS
C. SnS
D. Equal solubility for all

## D Watch Video Solution

12. In equalitative analysis, cations of graph $I I$ as well as group $I V$ both are precipitated in the form of sulphides. Due to low value of $K_{s p}$ of group II sulphides, group reagent is $\mathrm{H}_{2} \mathrm{~S}$ in the presence of dil. $\mathrm{HC1}$, and due to high value of $K_{s p}$ of group $I V$ sulphides, group reagent is $H_{2} S$ in the presence of $\mathrm{NH}_{4} \mathrm{OH}$ and $\mathrm{NH}_{4} \mathrm{C}$. In a solution containing 0.1 M each of $\mathrm{Sn}^{2+}, \mathrm{Cd}^{2+}$, and $\mathrm{Ni}^{2+}$ ions, $\mathrm{H}_{2}$ Sgas is passed.
$K_{s p} o f S n S=8 \times 10^{-29}, K_{s p} o f C d S=10^{-28}, K_{s p} o f N i S-3 \times 10^{-21}$
$K_{1} o f H_{2} S=1 \times 10^{-7}, K_{2} o f H_{2} S=1 \times 10^{-14}$
If $0.1 M H C 1$ is mixed in the solution containing only $0.1 M C d^{2+}$ ions and saturated with $H_{2} S$, then $\left[C d^{2+}\right]$ remaining in the solution after $C d S$ stopes to precipitate is:
A. $10^{-8}$
B. $8.2 \times 10^{-9}$
C. $5.6 \times 10^{-6}$
D. $5.6 \times 10^{-10}$

## Answer: A

## D Watch Video Solution

13. The degree of dissociation of weak electrolyde is inversely proportional to the square root fo concentration. It is called Ostwald's dilution law.
$\alpha=\sqrt{\frac{K_{a}}{c}}$ As the tempertaure increases, degree of dissociation will increase.
$\frac{\alpha_{1}}{\alpha_{2}}=\sqrt{\frac{K_{a_{1}}}{K_{a_{2}}}}$ if concentration is same.
$\frac{\alpha_{1}}{\alpha_{2}}=\sqrt{\frac{c_{2}}{c_{1}}}$ if acid is same.
$0.01 \mathrm{MCH}_{3} \mathrm{COOH}$ has $4.24 \%$ degree of dissociation, the degree of dissociation of $0.1 \mathrm{MCH}_{3} \mathrm{COOH}$ will be
A. $1.33 \%$
B. 4.24 \%
C. 5.24 \%
D. 0.33 \%

## Answer: A

## - Watch Video Solution

14. The degree of dissociation of weak electrolyde is inversely proportional to the square root fo concentration. It is called Ostwald's dilution law.
$\alpha=\sqrt{\frac{K_{a}}{c}}$ As the tempertaure increases, degree of dissociation will increase.
$\frac{\alpha_{1}}{\alpha_{2}}=\sqrt{\frac{K_{a_{1}}}{K_{a_{2}}}}$ if concentration is same.
$\frac{\alpha_{1}}{\alpha_{2}}=\sqrt{\frac{c_{2}}{c_{1}}}$ if acid is same.
$p H$ of $0.005 \mathrm{MHCOOH}\left[K_{a}=2 \times 10^{-4}\right]$ is equal to
A. 3
B. 2
C. 4
D. 5

## Answer: A

## - Watch Video Solution

15. The degree of dissociation of weak electrolyde is inversely proportional to the square root fo concentration. It is called Ostwald's dilution law.
$\alpha=\sqrt{\frac{K_{a}}{c}}$ As the tempertaure increases, degree of dissociation will increase.
$\frac{\alpha_{1}}{\alpha_{2}}=\sqrt{\frac{K_{a_{1}}}{K_{a_{2}}}}$ if concentration is same.
$\frac{\alpha_{1}}{\alpha_{2}}=\sqrt{\frac{c_{2}}{c_{1}}}$ if acid is same.
$a_{1}$ and $a_{2}$ are in ratio of $1: 2, K_{a_{1}}=2 \times 10^{-4}$. What will be $K_{a_{2}}$ ?
A. $8 \times 10^{-4}$
B. $2 \times 10^{-4}$
C. $4 \times 10^{-4}$
D. $1 \times 10^{-4}$

## Answer: A

## D Watch Video Solution

16. The following solutions are mixed: $500 \mathrm{mLof} 0.01 \mathrm{MAgNO}_{3}$ and 500 mL solution that was both 0.01 M in NaCI and 0.01 M in NaBr . Given $K_{\text {sp }} A G C I=10^{-10}, K_{s p} A g B r=5 \times 10^{-13}$.

Calculate the $\left[C I^{\Theta}\right]$ in the equilibrium solution.
A. $5 \times 10^{-5} M$
B. $2.5 \times 10^{-5}$
C. $5 \times 10^{-3} M$
D. $2.5 \times 10^{-3} \mathrm{M}$

## Answer: C

## - Watch Video Solution

17. The following solutions are mixed: $500 \mathrm{mLof} 0.01 \mathrm{MAgNO}_{3}$ and 500 mL solution that was both 0.01 M in NaCI and 0.01 M in NaBr . Given $K_{\text {sp }} A G C I=10^{-10}, K_{s p} A g B r=5 \times 10^{-13}$.

Calculate the $\left[A g^{\oplus}\right]$ in the equilibrium solution.
A. $2.0 \times 10^{-8} \mathrm{M}$
B. $2.0 \times 10^{-10} M$
C. $2.5 \times 10^{-5} \mathrm{M}$
D. $2.5 \times 10^{-8} \mathrm{M}$

## - Watch Video Solution

18. The following solutions are mixed: $500 \mathrm{mLof} 0.01 \mathrm{MAgNO}_{3}$ and 500 mL solution that was both 0.01 M in NaCI and 0.01 M in NaBr . Given $K_{s p} A G C I=10^{-10}, K_{s p} A g B r=5 \times 10^{-13},\left[A g^{+}\right]=2 \times 10^{-8}$.

Calculate the $\left[B r^{\Theta}\right]$ in the equilibrium solution.
A. $2.0 \times 10^{-8} M$
B. $2.0 \times 10^{-10} M$
C. $2.5 \times 10^{-5} M$
D. $2.5 \times 10^{-8} M$

## Answer: C

## - Watch Video Solution

19. When 1.5 mol of $\mathrm{CuCI}_{2} \cdot 2 \mathrm{H}_{2} \mathrm{O}$ is dissolved in enough water to make 1.0 L of solution.

Given: $K_{f} C u C I^{\Theta} 1.0\left(K_{f}\right.$ is the formation constant of $\left.\mathrm{CuCi}^{\oplus}\right)$ $\left[\mathrm{Cu}^{2+}\right]$ in solution is
A. 1.0 M
B. 0.5 M
C. 2.0 M
D. None

## Answer: A

## - Watch Video Solution

20. When 1.5 mol of $\mathrm{CuCI}_{2} \cdot 2 \mathrm{H}_{2} \mathrm{O}$ is dissolved in enough water to make 1.0 L of solution.

Given: $K_{f} \mathrm{CuCI}^{\Theta} 1.0\left(K_{f}\right.$ is the formation constant of $\left.\mathrm{CuCi}^{\oplus}\right)$ $\left[C I^{\Theta}\right]$ in solution is
A. 2.0 M
B. 1.0 M
C. 3.0 M
D. None

## Answer: A

## - Watch Video Solution

21. When 1.5 mol of $\mathrm{CuCI}_{2} \cdot 2 \mathrm{H}_{2} \mathrm{O}$ is dissolved in enough water to make 1.0 L of solution.

Given: $K_{f} \mathrm{CuCI}^{\Theta} 1.0\left(K_{f}\right.$ is the formation constant of $\left.\mathrm{CuCi}^{\oplus}\right)$
$\left[\mathrm{CiCI}^{\oplus}\right]$ in solution is
A. 1.0 M
B. 2.0 M
C. 3.0 M
D. 0.5 M

## Answer: B

## - Watch Video Solution

22. Acid rain takes place dur to combination of acidic oxides with water and it is an environmental concern all over the world. Assuming rain water is uncontaminated with $\mathrm{HNO}_{3}$ or $\mathrm{H}_{2} \mathrm{SO}_{4}$ and is in equilibrium with $1.25 \times 10^{-4} \mathrm{atmCO} 2$. The Henry's law constant $\left(K_{H}\right)$ is $1.25 \times 10^{6}$ torr. $K_{a_{1}}$ of $\mathrm{H}_{2} \mathrm{CO}_{3}=4.3 \times 10^{-7}$

What is the pH of neutral rain water ?
A. 5.64
B. 7.00
C. 5.85
D. 7.40

## Answer: C

23. Acid rain takes place dur to combination of acidic oxides with water and it is an envirronmental concern all over the world. Assuming rain water is uncontaminated with $\mathrm{HNO}_{3}$ or $\mathrm{H}_{2} \mathrm{SO}_{4}$ and is in equilibrium with $1.25 \times 10^{-4} \mathrm{~atm} \mathrm{CO}_{2}$. The Henry's law constant $\left(K_{H}\right)$ is $1.25 \times 10^{6}$ torr. $K_{a_{1}}$ of $\mathrm{H}_{2} \mathrm{CO}_{3}=4.3 \times 10^{-7}$

Given : $K_{f} C u C I^{\Theta}=1.0\left(K_{f}\right.$ is formation constant of $\left.\mathrm{CuCI}^{\oplus}\right)$
If $\mathrm{SO}_{2}$ content is the atomsphere is 0.64 ppm by volume, pH of rain water is (assume $100 \%$ ionisation of acid rain as monobasic acid).
A. 4.0
B. 5.0
C. 6.0
D. 7.0

## Answer: B

## - Watch Video Solution

24. In atmosphere, $\mathrm{SO}_{2}$ and NO are oxidised to $\mathrm{SO}_{3}$ and $\mathrm{NO}_{2}$, respectively,w hcih react with water to given $\mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{HNO}_{3}$. The resultant solution is called acid rain. $\mathrm{SO}_{2}$ dissolves in water to form diprotic acid.
$\mathrm{SO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{HSO}_{3}^{\Theta}+H^{\oplus}, K_{a_{1}}=10^{-2}$.
$\mathrm{HSO}_{3}^{\Theta} \Leftrightarrow \mathrm{SO}_{3}^{2-}+H^{\oplus}, K_{a_{2}}=10^{-7}$
and for equilibrium,
$\mathrm{SO}_{2}(a q)+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{SO}_{3}^{2-}(a q)+2 \mathrm{H}^{\oplus}(a q)$
$K_{a}=K_{a_{1}} \times K_{a_{2}}=10^{-9} a t 300 \mathrm{~K}$.
Which of the following reagnets will given white precipitate with the aqueous solution of sulphurous acid?
A. $\mathrm{BaCI}_{2}$
B. HCI
C. NaCI
D. KCI

## Answer: A

25. In atmosphere, $\mathrm{SO}_{2}$ and NO are oxidised to $\mathrm{SO}_{3}$ and $\mathrm{NO}_{2}$, respectively,w hcih react with water to given $\mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{HNO}_{3}$. The resultant solution is called acid rain. $\mathrm{SO}_{2}$ dissolves in water to form diprotic acid.
$\mathrm{SO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{HSO}_{3}^{\Theta}+H^{\oplus}, K_{a_{1}}=10^{-2}$.
$\mathrm{HSO}_{3}^{\Theta} \Leftrightarrow \mathrm{SO}_{3}^{2-}+H^{\oplus}, K_{a_{2}}=10^{-7}$
and for equilibrium,
$\mathrm{SO}_{2}(a q)+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{SO}_{3}^{2-}(a q)+2 \mathrm{H}^{\oplus}(a q)$
$K_{a}=K_{a_{1}} \times K_{a_{2}}=10^{-9} a t 300 \mathrm{~K}$.
The pH of 0.01 M aqueous solutioon of sodium sulphite $\left(\mathrm{Na}_{2} \mathrm{SO}_{3}\right)$
A. 4.5
B. 8.5
C. 9.0
D. 9.5

## - Watch Video Solution

26. In atmosphere, $\mathrm{SO}_{2}$ and NO are oxidised to $\mathrm{SO}_{3}$ and $\mathrm{NO}_{2}$, respectively,w hcih react with water to given $\mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{HNO}_{3}$. The resultant solution is called acid rain. $\mathrm{SO}_{2}$ dissolves in water to form diprotic acid.
$\mathrm{SO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \Leftrightarrow \mathrm{HSO}_{3}^{\Theta}+H^{\oplus}, K_{a_{1}}=10^{-2}$.
$\mathrm{HSO}_{3}^{\Theta} \Leftrightarrow \mathrm{SO}_{3}^{2-}+H^{\oplus}, K_{a_{2}}=10^{-7}$
and for equilibrium,
$\mathrm{SO}_{2}(a q)+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{SO}_{3}^{2-}(a q)+2 \mathrm{H}^{\oplus}(a q)$
$K_{a}=K_{a_{1}} \times K_{a_{2}}=10^{-9} a t 300 \mathrm{~K}$.
The dominant equilibrium in an aqueous solution of sodium hydrogen sulphite $\left(\mathrm{NaHSO}_{3}\right)$ is
$2 \mathrm{HSO}_{3}^{\Theta}(a q) \Leftrightarrow \mathrm{SO}_{2}(a q)+\mathrm{SO}_{3}^{2-}(a q)+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
The equilibrium constant for the above reaction is
A. $10^{-3}$
B. $10^{-5}$
C. $10^{-6}$
D. $10^{-9}$

## Answer: B

## - Watch Video Solution

27. In atmosphere, $\mathrm{SO}_{2}$ and NO are oxidised to $\mathrm{SO}_{3}$ and $\mathrm{NO}_{2}$, respectively,w hcih react with water to given $\mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{HNO}_{3}$. The resultant solution is called acid rain. $\mathrm{SO}_{2}$ dissolves in water to form diprotic acid.
$\mathrm{SO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{HSO}_{3}^{\Theta}+H^{\oplus}, K_{a_{1}}=10^{-2}$.
$\mathrm{HSO}_{3}^{\Theta} \Leftrightarrow \mathrm{SO}_{3}^{2-}+H^{\oplus}, K_{a_{2}}=10^{-7}$
and for equilibrium,
$\mathrm{SO}_{2}(a q)+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \Leftrightarrow \mathrm{SO}_{3}^{2-}(a q)+2 \mathrm{H}^{\oplus}(a q)$
$K_{a}=K_{a_{1}} \times K_{a_{2}}=10^{-9} a t 300 K$.
Which of the following statement is correct?
A. $\mathrm{H}_{2} \mathrm{SO}_{3}$ is less acidic than $\mathrm{H}_{2} \mathrm{SO}_{4}$.
B. $\mathrm{HNO}_{3}$ is less acidic than $\mathrm{HNO}_{2}$.
C. $\mathrm{SO}_{2}(g)$ is reduced in the atmosphere during thunderstron.
D. $\mathrm{CO}_{2}$ gas develop more acidity in rain water than $\mathrm{SO}_{2}$.

## Answer: A

## D Watch Video Solution

28. In acid-base titration react rapidly to neutralise each other. Equivalence point is a point at which the acid and the base (or oxidising agent and reducing agent) have beem added in equivalent quantities. The end point in the point at which the titration stops. since the purpose of the indicator is to stop the titration close to the point at which the acid and base were added in equivalent quantities, it is important that the equivalent point and the end point be as close as must change colour at a pH close to that of a solution of the salt of the acid base. Singificantly, the $p H$ changes most rapidly near the equivalent point. The exact shape of a titration curve depends on $K_{a}$ and $K_{b}$ of acid and base.


The following curve represents titration curve of HCI against KOH . The $p H$ at equivalent point is

Examine the titration curve below and answer the question.

A. 3
B. 6
C. 7
D. 8

## Answer: C

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29. In acid-base titration react rapidly to neutralise each other. Equivalence point is a point at which the acid and the base (or oxidising agent and reducing agent) have beem added in equivalent quantities. The end point in the point at which the titration stops. since the purpose of the indicator is to stop the titration close to the point at which the acid and base were added in equivalent quantities, it is important that the equivalent point and the end point be as close as must change colour at a $p H$ close to that of a solution of the salt of the acid base. Singificantly, the pH changes most rapidly near the equivalent point. The exact shape of a titration curve depends on $K_{a}$ and $K_{b}$ of acid and base.


The curve represents the titration of
A. CsOHbyHBr
B. HCIbyNaOH
C. HCl by ammonium hydroxide
D. $\mathrm{NH}_{3} \mathrm{byHNO} 3$

## Answer: A

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30. 



The suitable indicator for the titration is
A. Methy 1 orange
B. Bromothymol
C. Methy1 red
D. All of these

## Answer: D

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31. In acid-base titration react rapidly to neutralise each other. Equivalence point is a point at which the acid and the base (or oxidising
agent and reducing agent) have beem added in equivalent quantities. The end point in the point at which the titration stops. since the purpose of the indicator is to stop the titration close to the point at which the acid and base were added in equivalent quantities, it is important that the equivalent point and the end point be as close as must change colour at a $p H$ close to that of a solution of the salt of the acid base. Singificantly, the pH changes most rapidly near the equivalent point. The exact shape of a titration curve depends on $K_{a}$ and $K_{b}$ of acid and base.


The pH at equivalence point is
A. 2
B. 3
C. 7
D. 11

## Answer: C

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32. In acid-base titration react rapidly to neutralise each other. Equivalence point is a point at which the acid and the base (or oxidising agent and reducing agent) have beem added in equivalent quantities. The end point in the point at which the titration stops. since the purpose of the indicator is to stop the titration close to the point at which the acid and base were added in equivalent quantities, it is important that the equivalent point and the end point be as close as must change colour at a $p H$ close to that of a solution of the salt of the acid base. Singificantly, the $p H$ changes most rapidly near the equivalent point. The exact shape of a titration curve depends on $K_{a}$ and $K_{b}$ of acid and base.


Which of the following curves indicates the titration of a weak diprotic acid by KOH of equivalent strength?

A.

B.

C.


## D Watch Video Solution

33. Physical and chemical equilibrium can respond to a change in their pressure, temperature, and concentration of reactants and products. To describe the change in the equilibrium we have a principle named Le Chatelier principle. According to this principle, even if we make some changes in equilibrium, then also the system even re-establishes the equilibrium by undoing the effect.

In the reaction $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \Leftrightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$. If we increase the pressure of the system, the equilibrium is
A. Shifts in the product side
B. Remains unchanged
C. Shifts in the reactant side
D. Cannot be predicted

## Answer: A

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34. Physical and chemical equilibrium can respond to a change in their pressure, temperature, and concentration of reactants and products. To describe the change in the equilibrium we have a principle named Le Chatelier principle. According to this principle, even if we make some changes in equilibrium, then also the system even re-establishes the equilibrium by undoing the effect.

If we add $\mathrm{SO}_{4}^{2-}$ ion to a saturated solution of $\mathrm{Ag}_{2} \mathrm{SO}_{4}$, it will result in $a / / a n$
A. Result in an increase in $\mathrm{Ag}^{\oplus}$ concentration
B. Result in a decrease in $\mathrm{Ag}^{\oplus}$ concentration
C. Shift $\mathrm{Ag}^{\oplus}$ ions from solid $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ into solution.
D. Result in a decrease the $\mathrm{CrO}_{4}^{2-}$ ion concentration in the solution.

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35. Physical and chemical equilibria can respond to a change in their pressure, temperature, and concentration of reactants and products. To describe the change in the equilibrium, we have a principle named Le Chatelier's principle. This we can define in terms of energy, as the free energy change in equilibrium is zero means the system is stable. So if we are doing some changes in equilibrium, then the system having a tendency to reestablish the equilibrium by undoing the effect we brought. Consider the following equilibrium.

Three sparingly soluble salts $A_{2} B, A B$, and $A B_{3}$ are given. If all the three having the same value of solubility products $\left(K_{s p}\right)$, in the saturated solution, the correct order of their solubilites is
A. $A B_{3}>A B>A_{2} B$
B. $A B_{3}>A_{2} B>A B$
C. $A B>A B_{3}>A_{2} B$
D. $A B>A_{2} B>A B_{3}$

## Answer: D

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36. $\mathrm{H}_{3} \mathrm{PO}_{4}$ is a tribasic acid with $p K_{a_{1},} p K_{a_{2}}$ and $p K_{a_{3}} 1.12,7.21$, and 12.32 , respectively. It is used in fertiliser productions and its various salts are used in food, detrgent, toothpaste, and in metal treatment.

Small quantities of $\mathrm{H}_{3} \mathrm{PO}_{4}$ are used in inparting the sour or tart taste of soft drinkes, such as Coca Cola, and beers, in which $\mathrm{H}_{3} \mathrm{PO}_{4}$ is prese4nt $0.05 \%$ by weight (density $=1.0 \mathrm{gmL}^{-1}$ ).
$10^{-3} \mathrm{MH}_{3} \mathrm{PO}_{4}(\mathrm{pH}=7)$ is used in fertilisers as an aqueous soil digesting. Plants can absorb zinc in whater soluble from only. Zinc phosphate is the source of zinc and $\mathrm{PO}_{4}^{3-}$ ) ions in the soil. $K_{s p}$ of zinc phosphate $=9.1 \times 10^{-33}$.

Calculate the pH of a Coca Cola, assuming that the acidity of the cola arises only from $H_{3} \mathrm{PO}_{4}$ and $K_{a_{2}}$ and $K_{a_{3}}$ are no importance.
A. . 18
B. 2.2
C. 3.3
D. 4.4

## Answer: B

## - Watch Video Solution

37. $\mathrm{H}_{3} \mathrm{PO}_{4}$ is a tribasic acid with $p K_{a_{1}}, p K_{a_{2}}$ and $p K_{a_{3}} 1.12,7.21$, and 12.32, respectively. $10^{-3} \mathrm{MH}_{3} \mathrm{PO}_{4}(\mathrm{pH}=7)$ is used in fertilizers as an aqueous soil digesting. Plants can absorb zinc in water soluble from only. Zinc phosphate is the source of zinc and $\mathrm{PO}_{4}^{3-}$ ) ions in the soil. $K_{s p}$ of zinc phosphate $=9.1 \times 10^{-33}$.
$\left[\mathrm{PO}_{4}^{3-}\right]$ ion in the soil with $p H=7$, is
A. $10^{-3} \mathrm{M}$
B. $1.2 \times 10^{-4} \mathrm{M}$
C. $2.2 \times 10^{-4} \mathrm{M}$
D. $1.1 \times 10^{-10} M$

## Answer: C

## - Watch Video Solution

38. $\mathrm{H}_{3} \mathrm{PO}_{4}$ is a tribasic acid with $p K_{a_{1}}, p K_{a_{2}}$ and $p K_{a_{3}} 1.12,7.21$, and 12.32,
respectively. $10^{\wedge}(-3) \mathrm{M} \quad$ 'H_(3)PO_(4)'(pH $\quad=\quad$ 7)
. KspofZ $\in$ cphospê $=9.1 \times 10^{-33}$.
$\left[\mathrm{Zn}^{2+}\right]$ ion in the soil is
A. $2.9 \times 10^{-11} M$
B. $4.0 \times 10^{-10} M$
C. $3.0 \times 10^{-6} M$
D. $9.1 \times 10^{-5} M$

## Answer: A

39. The number of moles of $\mathrm{KMnO}_{4}$ required to oxidise 1 mol of $\mathrm{Fe}\left(\mathrm{C}_{2} \mathrm{O}_{4}\right)$ in acidic medium is
A. $6 \times 10^{-6}, 6 \times 10^{-6}$
B. $6 \times 10^{-6}, 1.5 \times 10^{-5}$
C. $1.5 \times 10^{-5}, 6 \times 10^{-6}$
D. $6 \times 10^{-6}, 3 \times 10^{-6}$

## Answer: B

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40. Aqueous solutions of $\mathrm{Na}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ and $\mathrm{CaCI}_{2}$ are mixed and precipitate of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ formed is filered and dried. 250 mL of the saturated solution of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ required 6.0 mL of $0.001 \mathrm{MKMnO}_{4}$ solution in acidic medium for complete titration.

Equivalent of $\mathrm{KMNO}_{4}$ required in the titration and equivalent of $\mathrm{C}_{2} \mathrm{O}_{4}^{2-}$ ion present in $\mathrm{CaC}_{2} \mathrm{O}_{4}$, respectively, are
A. $3 \times 10^{-5}, 3 \times 10^{-5}$
B. $1.8 \times 10^{-5}, 3 \times 10^{-6}$
C. $3 \times 10^{-6}, 6 \times 10^{-6}$
D. $6 \times 10^{-6}, 3 \times 10^{-6}$

## Answer: A

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41. Aqueous solutions of $\mathrm{Na}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ and $\mathrm{CaCI}_{2}$ are mixed and precipitate of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ formed is filered and dried. 250 mL of the saturated solution of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ required 6.0 mL of $0.001 \mathrm{MKMnO}_{4}$ solution in acidic medium for complete titration. $K_{s p}$ of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ is
A. $2.25 \times 10^{-12}$
B. $2.25 \times 10^{-10}$
C. $3.6 \times 10^{-9}$
D. $4.0 \times 10^{-9}$

## Answer: C

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42. Aqueous solutions of $\mathrm{Na}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ and $\mathrm{CaCI}_{2}$ are mixed and precipitate of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ formed is filered and dried. 250 mL of the saturated solution of $\mathrm{CaC}_{2} \mathrm{O}_{4}$ required 6.0 mL of $0.001 \mathrm{MKMnO}_{4}$ solution in acidic medium for complete titration.

Which is the indicator in the above titration?
A. Phenolphthalein
B. Methy1 ornage
C. $\mathrm{KMnO}_{4}$ it self
D. None

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43. Hin is an acidic indicator $\left(K_{\text {Ind }}=10^{-7}\right)$ which dissociates into aqueous acidic solution of 30 mL of $0.05 \mathrm{MH}_{3} \mathrm{PO}_{4}\left(K_{1}=10^{-3}, K_{2}=10^{-7}, K_{3}=10^{-13}\right)$

Calculate the $\left[\frac{\operatorname{Ind}^{\Theta}}{H \in}\right]$
A. $1.413 \times 10^{-5}$
B. $1.413 \times 10^{-4}$
C. $3.128 \times 10^{-5}$
D. $3.128 \times 10^{-14}$

## Answer: A

44. Hin is an acidic indicator $\left(K_{\text {Ind }}=10^{-7}\right)$ which dissociates into aqueous acidic solution of 30 mL of $0.05 \mathrm{MH}_{3} \mathrm{PO}_{4}\left(K_{1}=10^{-3}, K_{2}=10^{-7}, K_{3}=10^{-13}\right)$ If Hin and Ind ${ }^{\Theta}$ posses colour $P$ and $Q$, respectively, and concentration of HIn is 120 times than that of $\operatorname{Ind}^{\Theta}$. colour $Q$ predominates over $P$ when concnetration of $\operatorname{Ind}{ }^{\Theta}$ is 127 times of HIn.

What is the pH range of the indicator.
A. $4.896 \rightarrow 9.0792$
B. $4.896 \rightarrow 8.0792$
C. $4.896 \rightarrow 7.0792$
D. $4.896 \rightarrow 6.0792$

## Answer: A

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45. $H \in$ is an acidic indicator $\left(K_{\text {Ind }}=10^{-7}\right)$ which dissociates into aqueous acidic solution of 30 mL of $0.05 \mathrm{MH}_{3} \mathrm{PO}_{4}\left(K_{1}=10^{-3}, K_{2}=10^{-7}, K_{3}=10^{-13}\right)$

If this solution is treated with 30 mLof NaOH solution, then what molarity of NaOH is needed to reach the equivalence point with indicator?
A. 0.1 M
B. 0.2 M
C. 0.3 M
D. $0.4 M$

## Answer: A

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## Exercises Multiple Correct

1. 0.1 mol of $\mathrm{RNH}_{2}\left(K_{b}=5 \times 10^{-4}\right)$ is mixed with 0.08 mol of $\mathrm{HC1}$ and diluted to $1 L$. Calculate the $\left[H^{\oplus}\right]$ in the solution.
A. The concentration of $H^{\oplus}$ ion is $8 \times 10^{-11} M$.
B. The concentration of $H^{\oplus}$ ion is $8 \times 10^{-5} \mathrm{M}$.
C. The pH of solution is 9.8
D. The pOH of solution is 10.2 .

## Answer: A: C

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2. When weak base solution $\left(50 \mathrm{mLof0} 0.1 \mathrm{NNH}_{4} \mathrm{OH}\right)$ is titrated with strong acid ( 0.1 NHCD$)$, the pH of the solution initially decrease fast and then decreases slowely till near the equivalence point (as shown in figure).

Which of the following is//are correct.

A. The slow decrease of $p H$ is due to the formation of an acidic buffer solution after the addition of some HCI.
B. The slope of shown graph will be minimum when 25 mL of 0.1 NHCI is added.
C. The slow decrease of $p H$ is due to the formation of basic buffer solution.
D. The initial fast decrease in $p H$ is due to fast consumption of OH ions by HCI.

## Answer: B::C::D

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3. Which of the following statements about a weak acid strong base titration is//are correct?
A. The $p H$ after the equivalence point of the weal acid string base titration is determined by using the $K_{b}$ expression for the conjugate base.
B. A buffer solution of weal acid and its conjugate base is formed before the equivalence is reached.
C. The $p H$ at the equivalence point of a weak monoprotic acid strong base titration is equal to the $p H$ at the equivalence point of a strong acid-strong base titration.
D. The increase in $p H$ in the region near the equivalence point of a weak acid strong base titration is grater than the pH change in the same region of a strong acid strong base titration

## Answer: A:B

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4. An acid-base indicator has $K_{a}=10^{-5}$. The acid form of the indicator is red and basic form is blue. Which of the following is//are correct?
A. At $p H=4.52$, solution is red
B. At $\mathrm{pH}-5.47$, solution is blue.
C. At $p H=6$, solution is $75 \%$ red
D. At $p H=8$, solution is $75 \%$ blue.

## Answer: A::B

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5. When HCl gas is passed through a saturated solution of common salt, pure NaCl is Precipitated because:
A. HCl is higly soluble in water.
B. The ionic product $\left[N a^{\oplus}\right]\left[C l^{\Theta}\right]$ exceeds its solubility product $\left(K_{s p}\right)$.
C. The $K_{s p}$ of NaCl is lowered the presence of $\mathrm{HCl}^{\Theta}$ ions.
D. HCl causes precipitation.

## Answer: A::B::D

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6. Excess of $\mathrm{Ag}_{2} \mathrm{SO}_{4}(\mathrm{~s}), \mathrm{BaSO}_{4}(\mathrm{~s})$, and $\mathrm{Ba}_{3}\left(\mathrm{PO}_{4}\right)_{2}(s)$ are simultaneously in euqilibrium with distilled water. Which of the following is (are) true? Assume no hydrolysis of dissolved ions.
A. $\left[\mathrm{Ag}^{\oplus}\right]+2\left[\mathrm{Ba}^{2+}\right]=2\left[\mathrm{SO}_{4}^{2-}\right]+3\left[\mathrm{PO}_{4}^{3-}\right]$
B. $2\left[\mathrm{Ag}^{\oplus}\right]+4\left[\mathrm{Ba}^{2+}\right]=2\left[\mathrm{SO}_{4}^{2-}\right]+2\left[\mathrm{PO}_{4}^{3-}\right]$
C. $2\left[\mathrm{Ag}^{\oplus}\right]+3\left[\mathrm{Ba}^{2+}\right]=2\left[\mathrm{SO}_{4}^{2-}\right]+2\left[\mathrm{PO}_{4}^{3-}\right]$
D. $\left[\mathrm{Ag}^{\oplus}\right]+\left[\mathrm{Ba}^{2+}\right]=\left[\mathrm{SO}_{4}^{2-}\right]+\left[\mathrm{PO}_{4}^{3-}\right]$

## Answer: A

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7. $A$ solution is found to contain
$\left[C I^{\Theta}\right]=1.5 \times 10^{-1} M,\left[B r^{\Theta}\right]=5.0 \times 10^{-4} M,\left[\mathrm{CrO}_{4}^{2-}\right]=1.9 \times 10^{-2} M$.
A solution of $\mathrm{AgNO}_{3}$ (100 \% dissociated) is added to the above solution drop by drop. Which silver salt will precipiate first ? Given:

$$
K_{s p}(A g C I)=1.5 \times 10^{-10}, K_{s p}(\mathrm{AgBr})=5.0 \times 10^{-13}, K_{s p}\left(\mathrm{Ag}_{2} \mathrm{CrO}_{4}\right)=1.9 \times 10^{-12}
$$

A. $A g C I$
B. AgBr
C. $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$
D. $A g C I$ and $A g B r$ togther

## Answer: D

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8. $\mathrm{HgCrO}_{4}$ just begins to peripitate when equal volumes of $4 \times 10^{-4} \mathrm{MHg}_{2}\left(\mathrm{NO}_{3}\right)_{2}$ and $2 \times 10^{-5} \mathrm{MK}_{2} \mathrm{CrO}_{4}$ are combind. What is the approximate $K_{s p}$ value of $\mathrm{Hg}_{2} \mathrm{CrO}_{4}$ ?
A. $1 \times 10^{-18} \mathrm{molL}^{-1}$
B. $8 \times 10^{-9} \mathrm{molL}^{-1}$
C. $2 \times 10^{-9} \mathrm{molL}^{-1}$
D. $4 \times 10^{-9} \mathrm{molL}^{-1}$

## Answer: B

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9. What is general criteria of choosing a suitable indicator for a given titration?
A. The indicator should have a broad pH range.
B. $p H$ at the end point of titration should be close of neutral point of indicator
C. The indicator should have neutral point at $p H=7$.
D. The indicator must show a sharp colour changes near the equivalence point of titration point.

## Answer: B::D

10. Which of the following are true for an acid- base titration?
A. Indicators catalyse the acid-base reactions by releasing or accepting $H^{\oplus}$ ions.
B. Indicators do not significantly affect the pH of the solution to which they are added
C. Acid-base reactions do not occur in the absence of indicators
D. Indicators have different colours in dissociated and undissociated forms.

## Answer: B::D

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11. An acid-base indicator has $K_{a}=3.0 \times 10^{-5}$. The acid form of the indicator is red and the basic form is blue. Then:
A. $p H$ is 4.04 when indicator is $75 \%$ red.
B. $p H$ is 5.00 when indicator is $75 \%$ blue.
C. $p H$ is 5.00 when indicator is $75 \%$ red.
D. $p H$ is 5.05 when indicator is $75 \%$ blue.

## Answer: A::B

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12. At the end point, there is a sharp change of colour in the indicator. This happened because the
A. pH a end point changes sharply.
B. Structure of the indicator changes.
C. Colour of indicator is adsorbed by water.
D. Dissociation constant of acid and base differ by 10 .
13. For a series of indicators, the colour and pH range over which colour change takes place are as follows:

Indicator Colour change over pH range
$U \quad$ Yellow to blue pH 0.0 to 1.6
$V \quad$ Red to yellow pH 2.8 to 4.1
$W \quad$ Red to yellow pH 4.2 to 5.8
$X \quad$ Yellow to blue pH 6.0 to 7.7
$Y \quad$ Colourless to red pH 8.2 to 10.0
Which of the following statements is correct ?
A. Indicator $V$ could be used to find the equivalence point for 0.01 M acetic and 0.1 M ammonium hydroxide (ammonia solution) titration.
B. Indicator $Y$ could be used to distinguish between $0.1 M H C I$ and
0.01 MNaOH solutions in water.
C. Indicator $X$ could be used to distinguish between solution of ammonium chloride and sodium acetate.
D. Indicator $W$ could be suitable for use in determining the concentration of acetic acid in vinegar by base titration.

## Answer: C

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14. $H_{3} \mathrm{PO}_{4} \Leftrightarrow H^{\oplus}+H_{2} \mathrm{PO}_{4}^{\Theta}, K_{a_{1}}$ :
$\mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta} \Leftrightarrow H^{\oplus}+\mathrm{HPO}_{4}^{2-},{ }^{2} K_{a_{2}}:$
$\mathrm{HPO}_{4}^{2-} \Leftrightarrow H^{\oplus}+\mathrm{PO}_{4}^{3-}, K_{a_{3}}:$
Mark out the incorrect statements:
A. $K_{a_{1}}>K_{a_{2}}>K_{a_{3}}$
B. $p H\left(H_{2} P O_{4}^{\Theta}\right)=\frac{p K_{a_{1}}+p K_{a_{2}}}{2}$
C. Both $\mathrm{H}_{3} \mathrm{PO}_{4}$ and $\mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}$ are more acidic than $\mathrm{HPO}_{4}^{2-}$
D. Only $\mathrm{HPO}_{4}^{2-}$ is amphiprotic anion in the solution.

## Answer: B::D

15. Aqueous solution of $\mathrm{HNO}_{3}, \mathrm{CH}_{3}, \mathrm{CH}_{3} \mathrm{COOH}$, and $\mathrm{CH}_{3} \mathrm{COOK}$ of identical concentrations are given. The pair (s) of the solution which may form a buffer upon mixing is (are):
A. NaOH and $\mathrm{CH}_{3} \mathrm{COOH}$
B. $\mathrm{HNO}_{3}$ and $\mathrm{CH}_{3} \mathrm{COOK}$
C. $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COOK}$
D. $\mathrm{HNO}_{3}+\mathrm{CH}_{3} \mathrm{COOH}$

## Answer: A::B::C

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16. To which of the solution, addition of water would not effect the pH ? .
A. $100 \mathrm{mLof} 0.2 \mathrm{MCH}_{3} \mathrm{COOH}+100 \mathrm{mLof} 0.1 \mathrm{MNaOH}$
B. $100 \mathrm{mLof} 0.2 \mathrm{MCH}_{3} \mathrm{COOH}+100 \mathrm{mLof} 0.2 \mathrm{MNaOH}$
C. $200 \mathrm{mLof} 0.2 \mathrm{MCH}_{3} \mathrm{COOH}+100 \mathrm{mLof} 0.1 \mathrm{MNaOH}$
D. $100 \mathrm{mLof} 0.2 \mathrm{MCH}_{3} \mathrm{COOH}+200 \mathrm{mLof} 0.1 \mathrm{MNaOH}$

## Answer: A::C

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17. Which of the following salt solutions has $\mathrm{pH}<7$ ? .
A. $\mathrm{NH}_{4} \mathrm{~F}$
B. $\mathrm{Cr}\left(\mathrm{NO}_{3}\right)_{3}$
c. $\left[\left(\mathrm{CN}_{3}\right)_{3} \stackrel{\oplus}{N} H\right] C I^{\Theta}$
D. $\mathrm{CaI}_{2}$

## Answer: A::B::C

18. Which of the following represents hydrolysis?.
$\oplus$
A. $\mathrm{NH}_{4}+2 \mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{NH}_{4} \mathrm{OH}+\mathrm{H}_{3} \mathrm{O}^{\oplus}$
$\oplus$
B. $\mathrm{NH}_{4}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{NH}_{3}+\mathrm{H}_{3} \mathrm{O}^{\oplus}$
C. $\mathrm{HCO}_{3}^{\Theta}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{H}_{2} \mathrm{CO}_{3}+\stackrel{\ominus}{\mathrm{O}} \mathrm{H}$
D. $\mathrm{HCO}_{3}^{\Theta}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{CO}_{3}^{2-}+\mathrm{H}_{3} \mathrm{O}^{\oplus}$

## Answer: A:C

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19. The pH values of aqueous solutions of which of the following compounds does not change on dilution?
A. $\mathrm{PhCOONH}_{4}$
B. $\mathrm{NH}_{4} \mathrm{CN}$
C. HCOONa
D. $\mathrm{NH}_{4} \mathrm{CI}$

## Answer: A: B

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20. In $\mathrm{H}_{3} \mathrm{PO}_{4}$ which of the following is true?
A. $K_{a}=K_{a_{1}} \times K_{a_{2}} \times K_{a_{3}}$
B. $K_{a_{1}}<K_{a_{2}}<K_{a_{3}}$
C. $K_{a_{1}}>K_{a_{2}}>K_{a_{3}}$
D. $K_{a_{1}}=K_{a_{2}}=K_{a_{3}}$

## Answer: A:C

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21. Degree of hydrolysis ( $\alpha$ ) for a salt of strong acid and weak base is :
A. Is independent of dilution
B. Increases with dilution
C. Increases with decrease in $K_{b}$ of the base
D. Decreases with decrease in temperature

## Answer: B::C::D

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22. A solution containing a mixture of 0.05 MNaCI and 0.05 M Nal is taken. $\left(K_{s p} o f A g C I=10^{-10}\right.$ and $K_{s p}$ of $\left.\mathrm{AgI}=4 \times 10^{-16}\right)$. When $\mathrm{AgNO}_{3}$ is added to such a solution:
A. The concentartion fo $\mathrm{Ag}^{\oplus}$ required to precipitate $\mathrm{CI}^{\Theta}$ is

$$
2 \times 10^{-9} \mathrm{molL}^{-1}
$$

B. The concentartion of $\mathrm{Ag}^{\oplus}$ required to precipitate $I^{\Theta}$ is

$$
8 \times 10^{-15} \mathrm{molL}^{-1} .
$$

C. $A g C I$ and $A g I$ will be precipitate togther.
D. First AgI will be precipitated.

## Answer: A::B::D

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23. Which of the following is(are) correct when 0.1 L of $0.0015 \mathrm{MMgCl}_{2}$ and
0.1 L of 0.025 MNaF are mixed togther? $\left(K_{s p} o f M g F_{2}=3.7 \times 10^{-8}\right)$.
A. $M g F_{2}$ remains in solution
B. $M g F_{2}$ precipitates out
C. $\mathrm{MgCl}_{2}$ precipitates out
D. $\mathrm{Cl}^{\Theta}$ ions remains in solution

## Answer: B::D

24. Choose the correct statement:
A. pH of acidic buffer solution decrease if more salt is added
B. pH of acidic solution increases if more salt is added.
C. $p H$ of basic buffer increase if more salt is added.
D. $p H$ of basic buffer increase if more salt is added.

## Answer: B::C

## - Watch Video Solution

25. Which of the following is (are) correct for buffer solution?
A. Acidic buffer will be effective within in the $p H$ range $\left(p K_{a} \pm 1\right)$.
B. Basic buffer will be effective within the $p H$ range $\left(p K_{w}-p K_{b} \pm 1\right)$.
C. $\mathrm{H}_{3} \mathrm{PO}_{4}+\mathrm{NaH}_{2} \mathrm{PO}_{4}$ is not a buffer solution.
D. Buffers behave most effectively when the [Salt]/[Acid]

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26. A solution is prepared by dissolving 1.5 g of a monoacidic base into
1.5 kg of water at 300 K , which showed a depression in freezing point by $0.165^{\circ} \mathrm{C}$. When 0.496 g of the same base titrated, after dissolution, required 40 mL of semimolar $\mathrm{H}_{2} \mathrm{SO}_{4}$ solution. If $K_{f}$ of water is $1.86 \mathrm{Kkgmol}^{-1}$, then select the correct statements (s) out of the following( assuming molarity = molarity):
A. The pH of the solution of weak base is 12.9.
B. The ionisation constant of the base is $8 \times 10^{-3}$.
C. The osmotic pressure of the aqueous solution of base is 21.67 atm
D. The base is $10 \%$ ionized in aqueous solution.

## Answer: A::B::C::D

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27. A solution of $0.01 \mathrm{MFe}^{2+}$ in a saturated $\mathrm{H}_{2} \mathrm{~S}$ solution and (i) $0.2 \mathrm{MofH}^{\oplus}\left(\right.$ ii) $0.001 \mathrm{MofH}^{\oplus} .\left(K_{1} \times K_{2} o f H_{2} S=10^{-21}, K_{\text {sp }} \mathrm{FeS}=3.7 \times 10^{-19}\right)$
.Which of the following statements is//are correct
A. FeS will precipitate in solution (i).
B. FeS will not precipitate in solution (i).
C. FeS will precipitate in solution (ii).
D. FeS will precipitate in solution (ii).

## Answer: B::C

## - Watch Video Solution

28. Which statements is//are correct?
A. $0.1 \mathrm{MNH}_{3}$ solution will precipitate $\mathrm{Fe}(\mathrm{OH})_{2}$ from a 0.1 M solution $\mathrm{Fe}^{2+}$.
B. $0.1 \mathrm{MNH}_{3}$ solution will not precipitate $\mathrm{Mg}(\mathrm{OH})_{2}$ from a solution $\oplus$ which is 0.2 M in $\mathrm{NH}_{4}$ and 0.1 M in $\mathrm{Mg}^{2+}$
C. $0.1 \mathrm{MNH}_{3}$ solution will not precipitate AgOH from a solution which is $0.01 \mathrm{Min} \mathrm{Ag}^{\oplus}$.
D. Will precipitate is part (c).

## Answer: A::B::C

## - Watch Video Solution

29. Which statements is//are correct?
A. Compared to a strong acid, a weak acid titration with base starts at a higher pH .
B. Compared to a strong base, a weak base titration ends at a lower pH.
C. In both (a) and (b) titration curve is shortened at each end.
D. For titration of a weak base, the neraly vertical portion of the curve would be insufficient for an effective titration.

## Answer: A::B::C::D

## - Watch Video Solution

30. Which of the following solution will have $\mathrm{pH}=13$ ?
A. 2 gNaOHin 500 mL solution.
B. 100 mL solution fo $0.05 \mathrm{MCa}(\mathrm{OH})_{2}$.
C. 100 mL solution of $1.0 \mathrm{NCa}(\mathrm{OH})_{2}$.
D. 4 gNaOH in 500 mL solution.

## Answer: A::B::C

31. Which of the following statements (s) is (are) correct?
A. The pH of $1.0 \times 10^{-8} \mathrm{M}$ solution of HCI is 8 .
B. The conjugate base of $\mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}$ is $\mathrm{HPO}_{4}^{-2}$.
C. Autoprotoysis constant of water increases with temperature.
D. When a solution of weak monoprotic acid is titrated against a strong base, at half-neutralisation, point $p H=(1 / 2) p K_{a}$.

## Answer: B::C

## - Watch Video Solution

32. The pH of 0.1 M solution of the following salts decreases in the order
A. $\mathrm{NaCI}<\mathrm{NH}_{4} \mathrm{CI}<\mathrm{NaCN}<\mathrm{HCI}$
B. $\mathrm{HCI}<\mathrm{NH}_{4} \mathrm{CI}<\mathrm{NaCI}<\mathrm{NaCN}$
C. $\mathrm{NaCN}<\mathrm{NH}_{4} \mathrm{CI}<\mathrm{NaCI}<\mathrm{HCI}$
D. $\mathrm{HCI}<\mathrm{NaCI}<\mathrm{NaCN}<\mathrm{NH}_{4} \mathrm{CI}$

## Answer: B

## - Watch Video Solution

33. A buffer solution can be prepared from a mixture of
A. Sodium acetate and acetic acid in water.
B. Sodium acetat and hydrochloric acid in water.
C. Ammonia and ammonia chloride in water.
D. Ammonia and sodium hydroxide in water.

## Answer: A:C

## - Watch Video Solution

1. 100 mL of a buffer solution contains 0.1 M each of weak acid $H A$ and salt

NaA . How many gram of NaOH should be added to the buffer so that it $p H$ will be $6 ?\left(K_{a}\right.$ of $\left.H A=10^{-5}\right)$.
A. 0.328
B. 0458
C. 4.19
D. None

## Answer: A

## - Watch Video Solution

2. $\mathrm{K}_{a}$ for the reaction, $\mathrm{Fe}^{3+}($ aq. $)+\mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \Leftrightarrow \mathrm{Fe}(\mathrm{OH})^{2+}($ aq. $)+\mathrm{H}_{3} \mathrm{O}^{+}$(aq. $)$ is $6.5 \times 10^{-3}$ What is the max. pH value which could be used so that at least $80 \%$ of the total iron (III) in a dilute solution exists as $\mathrm{Fe}^{3+}$ ?
A. 2.0
B. $\sim 2.4$
C. $\sim 2.8$
D. $\sim 1.6$

## Answer: D

## - Watch Video Solution

3. The $p K_{b}$ of $C N^{\Theta}$ is 4.7. The $p H$ is solution prepared by mixing 2.5 mol of 2.5 mol of $K C N$ of 2.5 mol of $H C N$ in water and making the total volume upto 500 mL is
A. 10.3
B. 9.3
C. 8.3
D. 4.7

## Answer: B

4. A 0.1 molar solution of weak base BOH is $1 \%$ dissociated. If 0.2 mol of $B C l$ is added in $1 L$ solution of $B O H$. The degree of dissociation of $B O H$ will become
A. 0.02
B. 0.005
C. $5 \times 10^{-5}$
D. $2 \times 10^{-3}$

## Answer: C

## - Watch Video Solution

$$
\Theta
$$

5. If the equilibrium constant of $\mathrm{BOH} \leftrightarrow \mathrm{B}^{\oplus}+\mathrm{OH}$ at $25^{\circ} \mathrm{C}$ is $2.5 \times 10^{-6}$, then equilibrium constant for $\mathrm{BOH}+\mathrm{H}^{\oplus} \Leftrightarrow B^{\oplus}+\mathrm{H}_{2} \mathrm{O}$ at the same temperature is
A. $4.0 \times 10^{-9}$
B. $4.0 \times 10^{-5}$
C. $2.5 \times 10^{8}$
D. $2.5 \times 10^{-6}$

## Answer: C

## - Watch Video Solution

6. An aqueous solution of metal chloride $\mathrm{MCI}_{2}(0.05 M)$ is saturated with $\mathrm{H}_{2} \mathrm{~S}(0.1 \mathrm{M})$. The minimum pH at which metal sulphide will be precipiated is

$$
\left[K_{s p} M S=5 \times 10^{-21}, K_{1}\left(H_{2} S\right)=10^{-7}, K_{2}\left(H_{2} S\right)=10^{-14} .\right.
$$

A. 3.25
B. 2.50
C. 1.50
D. 1.25

## Answer: C

## D Watch Video Solution

7. The $p H$ of a solution of weak base at neutralisation with strong acid is
8. $K_{b}$ for the base is
A. $1.0 \times 10^{-4}$
B. $1.0 \times 10^{-6}$
C. $1.0 \times 10^{-8}$
D. None of these

## Answer: B

## - Watch Video Solution

8. The ionisation constant of an acid base indicator (a weak acid) is $1.0 \times 10^{-6}$. The ionised form of the indicator is red and unionised form is
blue. The pH change required to alter the colour of indicator from $80 \%$ red to $20 \%$ red is
A. 0.80
B. 1.20
C. 1.40
D. 2.00

## Answer: B

## - Watch Video Solution

9. $K_{s p}$ of $\mathrm{Mg}(\mathrm{OH})_{2}$ is $4.0 \times 10^{-6}$. At what minimum $\mathrm{pH}, \mathrm{Mg}^{2+}$ ions starts precipitating $0.01 \mathrm{MgCl}_{2}$
A. $2+\log 2$
B. $2-\log 2$
C. $12+\log 2$
D. $12-\log 2$

## Answer: C

## - Watch Video Solution

10. A solution of 0.1 MNaZ has $p H=8.90$. The $K_{a}$ of $H Z$ is
A. $6.3 \times 10^{-11}$
B. $6.3 \times 10^{-10}$
C. $1.6 \times 10^{-5}$
D. $1.6 \times 10^{-6}$

## Answer: C

## - Watch Video Solution

11. Phenolphalein does not act as an indicator for the titration between
A. HCI and $\mathrm{NH}_{4} \mathrm{OH}$
B. $\mathrm{Ca}(\mathrm{OH})_{2}$ and HCI
C. NaOH and $\mathrm{H}_{2} \mathrm{SO}_{4}$
D. KOH and $\mathrm{CH}_{3} \mathrm{COOH}$

## Answer: A

## D Watch Video Solution

12. The pink colour of phenolphthalein in alkaline medium is due to
$\Theta$
A. OH ions
B. Positive ion
C. Negative ion
D. Neutral form

## Answer: C

13. Methyl` orange gives red colour in
A. KOH solution
B. HCl solution
C. $\mathrm{Na}_{2} \mathrm{CO}_{3}$ solution
D. NaCl solution

## Answer: B

## - Watch Video Solution

14. A solution containing $\mathrm{NH}_{4} \mathrm{CI}$ and $\mathrm{NH}_{4} \mathrm{OH}$ has $\left[\begin{array}{l}\Theta \\ \mathrm{OH}\end{array}\right]=10^{-6} \mathrm{molL}^{-1}$, which of the following hydroxides would be precipitated when this solution in added in equal volume to a solution containing 0.1 M of metal ions?
A. $\mathrm{Mg}(\mathrm{OH})_{2},\left(K_{s p}=3 \times 10^{-11}\right)$
B. $\mathrm{Fe}(\mathrm{OH})_{2}\left(K_{\text {sp }}=8 \times 10^{-16}\right)$
C. $\mathrm{Cd}(\mathrm{OH})_{2}\left(K_{s p}=8 \times 10^{-6}\right)$
D. $\mathrm{AgOH}\left(K_{s p}=5 \times 10^{-3}\right)$

## Answer: B

## - Watch Video Solution

15. If equal volumes of $\mathrm{BaCl}_{2}$ and NaF solutions are mixed, which of these combination will not give a precipitate? $\left(K_{s p} o f B a F_{2}=1.7 \times 10^{-7}\right)$.
A. $10^{-3} \mathrm{BaCl}_{2}$ and $2 \times 10^{-2} \mathrm{MNaF}$
B. $10^{-3} \mathrm{MBaCl}_{2}$ and $1.5 \times 10^{-2} \mathrm{MNaF}$
C. $1.5 \times 10^{-2} \mathrm{MBaCl}_{2}$ and $10^{-2} \mathrm{MNaF}$
D. $2 \times 10^{-2} \mathrm{MBaCl}_{2}$ and $2 \times 10^{-2} \mathrm{MNaF}$

## Answer: C

16. The solubility of solid silver chromate, $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$, is determined in three solvents $K_{s p}$ of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}=9 \times 10^{-12}$
I. pure water II. $0.1 \mathrm{AgNO}_{3}$
III. $0.1 \mathrm{MNa}_{2} \mathrm{CrO}_{4}$

Predict the relative solubility of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$ in the three solvents.
A. $I=I I=I I I$
B. I $<$ II $<$ III
C. $I I=I I I<I$
D. II $<$ III $<$ I

## Answer: D

- Watch Video Solution

17. The solubility products of $\mathrm{Al}(\mathrm{OH})_{3}$ and $\mathrm{Zn}(\mathrm{OH})_{2}$ are $8.5 \times 10^{-23}$ and $1.8 \times 10^{-14}$ respectively. If $\mathrm{NH}_{4} \mathrm{OH}$ is added to a solution containing $\mathrm{Al}^{3+}$ and $\mathrm{Zn}^{2+}$ ions, then substance precipitated first is:
A. $\mathrm{Al}(\mathrm{OH})_{3}$
B. $\mathrm{Zn}(\mathrm{OH})_{2}$
C. Both (a) and (b)
D. None of these

## Answer: A

## - Watch Video Solution

18. If $K_{s p}\left(\mathrm{PbSO}_{4}\right)=1.8 \times 10^{-8}$ and $K_{a}\left(\mathrm{HSO}_{4}^{\Theta}\right)=1.0 \times 10^{-2}$ the equilibrium constant for the reaction.
$\mathrm{PbSO}_{4}(s)+H^{\oplus}(a q) \Leftrightarrow \mathrm{HSO}_{4}^{\Theta}(a q)+\mathrm{Pb}^{2+}(a q)$ is
A. $1.8 \times 10^{-6}$
B. $1.8 \times 10^{-10}$
C. $2.8 \times 10^{-10}$
D. $1.0 \times 10^{-2}$

## Answer: A

## - Watch Video Solution

19. Which one of the following is true for any diprotic acid, $\mathrm{H}_{2} \mathrm{X}$ ?
A. $K_{a_{2}}>K_{a_{1}}$
B. $K_{a_{1}}>K_{a_{2}}$
C. $K_{a_{2}}=\frac{1}{K_{a_{1}}}$
D. $K_{a_{2}}=K_{a_{1}}$

## Answer: B

20. The $K_{s p}$ of $\mathrm{Mg}(\mathrm{OH})_{2}$ is $1 \times 10^{-12} \cdot 0.01 \mathrm{MMg}^{2+}$ will precipitate at the limiting pH of
A. (a) 8
B. (b) 9
C. (c) 10
D. (d) 12

## Answer: B

## - Watch Video Solution

21. The solubility products of $M A, M B, M C$ and $M D$ are $1.8 \times 10^{-10}, 4 \times 10^{-3}, 4 \times 10^{-8}$ and $6 \times 10^{-5}$ respectively. If a 0.01 M solution of $M X$ is added dropwise to a mixture containing $A^{-}, B^{-}, C^{-}$and $D^{-}$ions, then the one to be precipitated first will be:
A. $M A$
B. $M B$
C. $M C$
D. $M D$

## Answer: A

## - Watch Video Solution

22. A solution is saturated with respect to $\mathrm{SrCO}_{3}$ and $\mathrm{SrF}_{2}$. The $\left[\mathrm{CO}_{3}^{2-}\right]$ was found to be $1.2 \times 10^{-3} \mathrm{M}$. The concnetration of $F^{\Theta}$ in the solution would be

Given $K_{s p}$ of $\mathrm{SrCO}_{3}=7.0 \times 10^{-10} \mathrm{M}^{2}$,

$$
K_{s p}{\mathrm{of} S r F_{2}}=7.9 \times 10^{-10} \mathrm{M}^{3},
$$

A. $1.3 \times 10^{-3} \mathrm{M}$
B. $2.6 \times 10^{-2} \mathrm{M}$
C. $3.7 \times 10^{-2} \mathrm{M}$
D. $5.8 \times 10^{-7} \mathrm{M}$

## D Watch Video Solution

23. The number of $S^{2-}$ ions present in 1 L of $0.1 \mathrm{MH}_{2} S\left[K_{a}\left(\mathrm{H}_{2} \mathrm{~S}\right)=10^{-21}\right]$ solution having $\left[H^{\oplus}\right]=0.1 M$ is:
A. (a) $6.023 \times 10^{3}$
B. (b) $6.023 \times 10^{4}$
C. (c) $6.023 \times 10^{5}$
D. (d) $6.023 \times 10^{6}$

## Answer: A

## - Watch Video Solution

24. The solubility of AgI in NaI solutions is less than that in pure water because:
A. (a) AgI forms complex with NaI
B. (b) Of common ion effect
C. (c) Solubility product of AgI is less than that of NaI .
D. (d) The temperature of the solution decreases.

## Answer: B

## - Watch Video Solution

25. Three sparigly soluble salts $M_{2} X, M X$, and $M X_{3}$ have the same solubility product. Their solubilities will be in the order
A. $M X_{3}>M X>M_{2} X$
B. $M X_{3}>M_{2} X>M X$
C. $M X>M X_{3}>M_{2} X$
D. $M X>M_{2} X>M X_{3}$

## Answer: B

26. When 0.2 M solution of acetic acid is neutralised with 0.2 MNaOH in 500 mL of water, the pH of the resulting solution will be: $\left[p K_{a}\right.$ of acetic acid $=4.74]$
A. 12.67
B. 7.87
C. 8.87
D. 7

## Answer: C

## - Watch Video Solution

27. A weak acid $H X$ has the dissociation constant $1 \times 10^{-5} \mathrm{M}$. It forms a salt NaX on reaction with alkali. The percentage hydrolysis of 0.1 M solution of NaX is
A. $0.001 \%$
B. $0.01 \%$
C. 0.1 \%
D. $0.15 \%$

## Answer: B

## - Watch Video Solution

28. A certain buffer solution contains equal concentartion of $X^{\Theta}$ and $H X$. The $K_{b}$ for $X^{\Theta}$ is $10^{-10}$. The $p H$ of the buffer is
A. 4
B. 7
C. 10
D. 14
29. $K_{a}$ of $H A$ is $10^{-4}$. The equilibrium constant for its reaction with a strong base is
A. $1.0 \times 10^{-4}$
B. $1.0 \times 10^{-10}$
C. $1.0 \times 10^{10}$
D. $1.0 \times 10^{14}$

## Answer: C

## - Watch Video Solution

30. Auto-ionisation of liquid $\mathrm{NH}_{3}$ is
$2 \mathrm{NH}_{3} \Leftrightarrow \mathrm{NH}_{4}^{\oplus}+\mathrm{NH}_{2}^{\Theta}$
with $K_{N H_{3}}=\left[\mathrm{NH}_{4}^{\oplus}\right]\left[\mathrm{NH}_{2}^{\Theta}\right]=10^{-30}$ at $-50^{\circ} \mathrm{C}$ Number of amide ions
$\left(\mathrm{NH}_{2}^{\Theta}\right)$, present per $\mathrm{mm}^{3}$ of pure liquid $\mathrm{NH}_{3}$ is
A. (a) 602
B. (b) 301
C. (c) 200
D. (d) 100

## Answer: A

## - Watch Video Solution

31. A mixture of weak acid is 0.1 M in $\mathrm{HCOOH}\left(K_{a}=1.8 \times 10^{-4}\right)$ and $0.1 M$ in $\operatorname{HOCN}\left(K_{a}=3.1 \times 10^{-4}\right)$. Hence, $\left[\mathrm{H}_{3} \mathrm{O}^{\oplus}\right]$ is
A. (a) $7.0 \times 10^{-3} M$
B. (b) $4.1 \times 10^{-4} M$
C. (c) 0.20 M
D. (d) $4.1 \times 10^{-3} M$

## - Watch Video Solution

32. pH of solution made by mixing 50 mL of $0.2 \mathrm{MNH}_{4} \mathrm{CI}$ and 75 mL of 0.1 MNaOH is $\left[p K_{b} o f \mathrm{NH}_{3}(a q)=4.74 \cdot \log 3=0.47\right]$
A. 7.02
B. 13.0
C. 7.02
D. 9.73

## Answer: D

## - Watch Video Solution

33. Some chemists at wished to perpare a saturated solution of a silver compound and they wanted it to have the highest concentration of silver ion possible. Which of the following compound would they use ?
$K_{s p}(A g C I)=1.8 \times 10^{-10}, K_{s p}(A g B r)=5.0 \times 10^{-13}$,
$K_{s p}\left(\mathrm{Ag}_{2} \mathrm{CrO}_{4}\right)=2.4 \times 10^{-12}[\mathrm{Use} 3 \sqrt{0.6}=0.84]$
A. AgCI
B. AgBr
C. $\mathrm{Ag}_{2} \mathrm{CrO}_{4}$
D. all of these

## Answer: C

## - Watch Video Solution

34. An acid-base indicator has a $K_{a}=3.0 \times 10^{-5}$. The acid form of the indicator is red and the basic form is blue. Then
A. $p H$ is 4.05 when indicator is $75 \%$ red.
B. $p H$ is 5.00 when indicator is $75 \%$ blue.
C. Both (a) and (b) are correct.
D. None of these

## Answer: C

## - Watch Video Solution

35. The $p H$ value of 0.001 M aqueous solution of NaCl is
A. (a) 7
B. (b) 4
C. (c) 11
D. (d) Unpredictable

## Answer: A

## - Watch Video Solution

36. Which of the following will supress the ionisation of acetic acid in aquoeus solution?
A. NaCI
B. HCI
C. $K C I$
D. Unpredictable

## Answer: B

## - Watch Video Solution

37. An aqueous solution of HCl is $10^{-9} \mathrm{MHCl}$. The pH of the solution should be
A. 9
B. Between 6 and 7
C. 7
D. Unpredictable

## Answer: B

## - Watch Video Solution

38. Which of the following represents the conjugate pair of $\mathrm{NH}_{3}$ ?
A. $\mathrm{NH}_{2}^{\Theta}$
B. $N H_{4}^{\oplus}$
C. Both (a) and (b)
D. $N^{3-}$

## Answer: C

## - Watch Video Solution

39. One of the following is a Bronsted acid but not a Bronsted base:
A. $\mathrm{H}_{2} \mathrm{~S}$
B. $\mathrm{H}_{2} \mathrm{~S}$
C. $\mathrm{HCO}_{3}{ }^{\Theta}$
D. $\mathrm{NH}_{3}$

## Answer: A

## - Watch Video Solution

40. In the third group of qualitive analysis, the precipitating reagent is $\mathrm{NH}_{4} \mathrm{CI} / \mathrm{NH}_{4} \mathrm{OH}$. The function of $\mathrm{NH}_{4} \mathrm{CI}$ is to
A. increases the ionisation of $\mathrm{NH}_{4} \mathrm{OH}$.
B. Supress the ionisation of $\mathrm{NH}_{4} \mathrm{OH}$.
C. Convert the ions of group theird into their respective chlorides.
D. Stabilise the hydroxides of group III cations.
41. At a certain temperature the value of $p K_{w}$ is 13.4 and the measured $p H$ of soln is 7. The solution is
A. Acidic
B. Basic
C. Neutral
D. Unpredictable

## Answer: B

## - Watch Video Solution

42. When 2 mol of HCI is added to 1 L of an acidic buffer, its pH changes from 3.4 to 2.9. The buffer capacity of the buffer solution is
A. 2
B. 0
C. 4
D. 8

## Answer: C

## - Watch Video Solution

43. Let the solubilities of AgCI in $\mathrm{H}_{2} \mathrm{O}$, and in $0.01 \mathrm{MCaCI}_{2}, 0.01 \mathrm{MNaCI}$, and $0.05 \mathrm{MAgNO}_{3}$ be $S_{1}, S_{2}, S_{3}, S_{4}$, respectively. What is the correct relationship between these quantites.
A. $S_{1}>S_{2}>S_{3}>S_{4}$
B. $S_{1}>S_{2}=S_{3}>S_{4}$
C. $S_{1}>S_{3}>S_{2}>S_{4}$
D. $S_{4}>S_{2}>S_{3}>S_{1}$

## Answer: C

44. Which of the following salts will not undergo hydrolysis in water?
A. Sodium sulphate
B. Ammonium sulphate
C. Aluminimum sulphate
D. All the salts will hydrolyse

## Answer: A

## - Watch Video Solution

45. Which of the following salts will not change the pH of pure water on dissociation?
A. (a) KCl
B. (b) $\mathrm{AICl}_{3}$
C. (c) $\mathrm{Na}_{2} \mathrm{CO}_{3}$
D. (d) $\mathrm{AI}_{2}\left(\mathrm{SO}_{4}\right)_{3}$

## Answer: A

## D Watch Video Solution

46. A salt $X$ is dissolved in water having $p H=7$. The resulting solution has a pH more than 7 . The salt is made by neutralisation of
A. A strong acid and strong base
B. A strong acid and strong weak base
C. A weak acid and weak base
D. A weak acid and strong base

## Answer: D

## - Watch Video Solution

47. The $p H$ of a solution 7.00 . To this solution, sufficient base is added to $\Theta$
increase the pH to 12.0. The increase in OH ion concentration is
A. $5 \times$
B. $100 \times$
C. $10^{5} \times$
D. $4 \times$

## Answer: C

## - Watch Video Solution

48. Assuming $\mathrm{H}_{2} \mathrm{SO}_{4}$ to be completely ionised the pH of a 0.05 M aqueous of sulphuric acid is approximately
A. 0.01
B. 0.005
C. 2

## D. 1

## Answer: D

## - Watch Video Solution

49. A solution has $p O H$ equal to 13 at $298 K$. The solution will be
A. Highly acidic
B. Highly basic
C. Moderatly basic
D. Unpredictable

## Answer: A

## - Watch Video Solution

50. If ammonia is added to pure water, the concentration of a chemical species already present will decrease. The species is
A. $O_{2}^{\Theta}$
$\Theta$
B. OH
C. $\mathrm{H}_{3} \mathrm{O}^{\oplus}$
D. $\mathrm{H}_{2} \mathrm{O}$

## Answer: C

## - Watch Video Solution

51. The pH of a dilute solution of acetic acid was found to be 4.3. The addition of a small crystal of sodium acetate will cause $p H$ to
A. (a) Become less than 4.3
B. (b) Become more than 4.3
C. (c) Remain equal to 4.3
D. (d) Unpredictable

## Answer: B

## - Watch Video Solution

52. Which of the following can act both as a Bronsted acid and a Bronsted base?
A. (a) $O_{2}^{\Theta}$
B. (b) HCl
C. (c) $\mathrm{HSO}_{4}^{\Theta}$
D. (d) $\mathrm{Na}_{2} \mathrm{CO}_{3}$

## Answer: C

## - Watch Video Solution

53. Which of the following is a Lewis base?
A. $\mathrm{H}_{2} \mathrm{O}$
B. $C I^{\Theta}$
C. $B F_{3}$
D. $\mathrm{NH}_{3}$

## Answer: C

## - Watch Video Solution

54. Which of the following is not a Lewis base?
A. $C N^{\Theta}$
B. ROH
C. $\mathrm{NH}_{3}$
D. $\mathrm{AICI}_{3}$

## - Watch Video Solution

## $\Theta$

55. Conjugate base of $O H$ is
A. $\mathrm{H}_{2} \mathrm{O}$
B. $\mathrm{H}_{3} \mathrm{O}^{\oplus}$
C. $H^{\oplus}$
D. $\mathrm{O}^{2-}$

## Answer: D

Watch Video Solution
56. Which of the following will have the largest pH ?
A. (a) $M / 10 H C I$
B. (b) $M / 100 H C I$
C. (c) $M / 10 \mathrm{NaOH}$
D. (d) $M / 100 \mathrm{NaOH}$

## Answer: C

## - Watch Video Solution

57. Which one of following will have the largest pH ?
A. Solution containing $1 \times 10^{-2}$ molofK $_{2} \mathrm{SO}_{4} L^{-1}$.
B. Pure water.
C. Solution containing $1.0 \times 10^{-2}$ molof CIL $^{-1}$.
D. Solution containing $1 \times 10^{2}$ molof $^{2} \mathrm{H}_{4} \mathrm{OHL}^{-1}$.

## Answer: D

58. When 20 mL of $\mathrm{M} / 20 \mathrm{NaOH}$ is added to 10 mL of $\mathrm{M} / 10 \mathrm{HCI}$, the resulting solution will
A. Turn blue litmus red.
B. Turn phenolpthalein solution pink.
C. Turns methy orange red.
D. Will have no effect on either red or blue litmus

## Answer: D

## - Watch Video Solution

59. pOH water is 7.0 at 298 K . If water is heated to 350 K , which of the following should be ture?
A. pOH will decrease
B. pOH will increase
C. pOH will remain seven
D. Concentration of $H^{\oplus}$ ions will increae but that of OH will decrease.

## Answer: A

## - Watch Video Solution

60. Solubility of salt $A_{2} B_{3}$ is $1 \times 10^{-4}$, its solubility product is
A. $1.08 \times 10^{20}$
B. $1.08 \times 10^{18}$
C. $2.6 \times 10^{-18}$
D. $1.08 \times 10^{-18}$

## Answer: D

## - Watch Video Solution

61. The value of $K_{s p}$ is $\mathrm{HgCl}_{2}$ at room temperature is $4.0 \times 10^{-15}$. The concentration of $\mathrm{Cl}^{-}$ion in its aqueous solution at saturation point is
A. $1 \times 10^{-5}$
B. $2 \times 10^{-5}$
C. $2 \times 10^{-15}$
D. $8 \times 10^{-15}$

## Answer: B

## - Watch Video Solution

62. At $90^{\circ} \mathrm{C}$, pure water has $\left[\mathrm{H}_{3} \mathrm{O}^{\oplus}\right]=10^{-7} \mathrm{molL}^{-1}$. What is the value of $K_{w}$ at $90^{\circ} \mathrm{C}$ ?
A. $10^{-6}$
B. $10^{-12}$
C. $10^{-13}$
D. $10^{-14}$

## Answer: C

## - Watch Video Solution

63. What is the solubility of $\mathrm{PbSO}_{4}$ in $0.01 \mathrm{MNa}_{2} \mathrm{SO}_{4}$ solution if $K_{s p}$ for $\mathrm{PbSO}_{4}=1.25 \times 10^{-9} ?$
A. $1.25 \times 10^{-7} \mathrm{molL}^{-1}$
B. $1.25 \times 10^{-9} \mathrm{molL}^{-1}$
C. $1.25 \times 10^{-10} \mathrm{molL}^{-1}$
D. $0.10 \mathrm{molL}^{-1}$

## Answer: A

64. The pH of an aqueous solution of $\mathrm{Ba}(\mathrm{OH})_{2}$ is 10 . If the $\mathrm{K}_{\text {sp }}$ of $\mathrm{Ba}(\mathrm{OH})_{2}$ is $1 \times 10^{-9}$, then the concentration of $\mathrm{Ba}^{2+}$ ions in the solution in $\mathrm{molL}^{-1}$ is
A. $1 \times 10^{-2}$
B. $1 \times 10^{-4}$
C. $1 \times 10^{-1}$
D. $1 \times 10^{-5}$

## Answer: C

## - Watch Video Solution

65. How many grams of NaOH must be dissolved in 1 L of the solution to give it a pH value of 12 ?
A. $0.20 g L^{-1}$
B. $0.40 \mathrm{gL}{ }^{-1}$
C. $0.10 g L^{-1}$
D. $1.2 g L^{-1}$

## Answer: B

## - Watch Video Solution

66. Which of the following solutions will have $\mathrm{pH}=10$ at 298 K ?
A. $1 \times 10^{-10} \mathrm{MHCI}$ solution
B. $1 \times 10^{-4} \mathrm{MNaOH}$ solution
C. $1 \times 10^{-10} \mathrm{MNaOH}$ solution
D. Both (a) and (b)

## Answer: B

## - Watch Video Solution

67. An acid $H A$ is $40 \%$ dissociated in an aqueous solution. The hydronium ion concentration of its 0.2 M solution would be
A. 0.08 M
B. $0.4 M$
C. 0.2 M
D. None

## Answer: A

## - Watch Video Solution

68. $20 \mathrm{~cm}^{3}$ of $x M$ solution of HCl is exactly neutralised by $40 \mathrm{~cm}^{3}$ of 0.05 MNaOH solutions, the pH of HCl solution is
A. 1.0
B. 2
C. 1.5
D. 2.5

## Answer: A

## - Watch Video Solution

69. A monoprotic acid $(H A)$ is $1 \%$ ionised in its aqueous solution of $0.1 M$ strength. Its pOH will be
A. 11
B. 3
C. 10
D. 2

## Answer: A

70. The pH of a solution is 5 . to this solution acid was added so that its pH value bcomes 2.0. The increase in $\mathrm{H}^{+}$concentration is:
A. 1000 times
B. 2.5 times
C. 100 times
D. 5 times

## Answer: A

## - Watch Video Solution

71. What would be the solubility of silver chloride in 0.10 MNaCI solution? $K_{\text {sp }} f$ or $A g C I=1.20 \times 10^{-10}$
A. 0.1 M
B. $1.2 \times 10^{-6} \mathrm{M}$
C. $1.2 \times 10^{-9} \mathrm{M}$
D. $1.2 \times 10^{-10} M$

## Answer: C

## - Watch Video Solution

72. Which of the following metal sulphides has maximum solubility in water?
A. (a) $\operatorname{CdS}\left(K_{s p}=36 \times 10^{-30}\right)$
B. (b) $\operatorname{FeS}\left(K_{\text {sp }}=11 \times 10^{-20}\right)$
C. (c) $\operatorname{HgS}\left(K_{\text {sp }}=32 \times 10^{-54}\right)$
D. (d) $\operatorname{ZnS}\left(K_{\text {sp }}=11 \times 10^{-22}\right)$

## Answer: B

## - Watch Video Solution

73. $M_{2} \mathrm{SO}_{4}\left(M^{\oplus}\right.$ is a monovalent metal ion) has a $K_{\text {sp }}$ of $3.2 \times 10^{-6}$ at 298 K . The maximum concentration of $\mathrm{SO}_{4}^{2-}$ ion that could be attained in a saturated solution of this solid at 298 K is
A. $3 \times 10^{-3} M$
B. $7 \times 10^{-2} M$
C. $2.89 \times 10^{-4} \mathrm{M}$
D. $2 \times 10^{-2} \mathrm{M}$

## Answer: D

## - Watch Video Solution

74. $K_{s p}$ for lead iodate $\left[\mathrm{Pb}\left(\mathrm{IO}_{3}\right)_{2}\right.$ is3.2 $\times 10^{-14}$ at a given temperature. The solubility in $\mathrm{molL}^{-1}$ will be
A. $2.0 \times 10^{-5}$
B. $\left(3.2 \times 10^{-7}\right)^{1 / 2}$
C. $\left(3.8 \times 10^{-7}\right)$
D. $4.0 \times 10^{-6}$

## Answer: A

## - Watch Video Solution

75. The pH of a 0.1 M solution of $\mathrm{NH}_{4} \mathrm{OH}$ (having dissociation constant $\left.K_{b}=1.0 \times 10^{-5}\right)$ is equal to
A. 10
B. 6
C. 11
D. 12

## Answer: C

76. The best indicator for the detection of end point in titration of a weak acid and a strong base is :
A. Methy1 orange ( pH range $3 \rightarrow 4$ )
B. Methy1 red ( pH range $4 \rightarrow 6$ )
C. Thymol blue ( pH range $8 \rightarrow 3$ )
D. Phemolphethalein ( pH range $8 \rightarrow 10$ )

## Answer: D

## - Watch Video Solution

77. When a solid KCI is added to a saturated solution of $\mathrm{AgClin}_{2} \mathrm{O}$,
A. Nothing happens.
B. Solubility of AgCI decreases.
C. Solubility of AgCI increases.
D. Solubility product of AgCI increases.

## Answer: B

## D Watch Video Solution

78. Two buffer solutions $A$ and $B$ each made with benzoic acid and sodium benzoate differ in their $p H$ by two units. A has salt: acida:b. $B$ has salt: acid $=b: a$. If $a>b$, then the value of $a: b$ is
a. $3.17 \quad$ b.10.0 $\quad$ c.3.0 $\quad$ d. 6.0
A. 10,000
B. 3.17
C. 6.61
D. 2.10

## Answer: B

79. $\mathrm{CaCO}_{3}$ and $\mathrm{BaCO}_{3}$ have solubility product values $1 \times 10^{-8}$ and $5 \times 10^{-9}$, respectively. If water is shaken up with both solids till equilibrium is reached, the concentration of $\mathrm{CO}_{3}^{2-}$ ion is
A. $1.5 \times 10^{-8}$
B. $1.225 \times 10^{-4}$
C. $2.25 \times 10^{-9}$
D. None of these

## Answer: B

## - Watch Video Solution

80. The $p H$ of an acidic buffer can be raised by 2 units by
A. Increasing the concentration of both weak acid and salt by two moles
B. Increasing the concentration of both the acid and salt by 10 times.
C. Diluting the solution by 10 times.
D. Increasing the concentration of the salt by 10 times by decreasing concentration of the acid by 10 times.

## Answer: D

## - Watch Video Solution

81. A buffer solution can be prepared from a mixture of
A. HCI and NaCI
B. $\mathrm{NaH}_{2} \mathrm{PO}_{4}$ and $\mathrm{Na}_{2} \mathrm{HPO}_{4}$
C. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{NaCI}$
D. $\mathrm{NH}_{4} \mathrm{OH}+\mathrm{NH}_{3}$

## Answer: B

82. 20 mol of $\mathrm{M} / 10 \mathrm{CH}_{3} \mathrm{COOH}$ solution is titrated with $\mathrm{M} / 10 \mathrm{NaOH}$ solution. After addition of 16 mL solution of NaOH . What is the pH of the solution $\left(p K_{a}=4.74\right)$
A. 5.05
B. 4.15
C. 4.75
D. 5.35

## Answer: D

## - Watch Video Solution

83. The $K_{s p}$ value of $\mathrm{CaCO}_{3}$ and $\mathrm{CaC}_{2} \mathrm{O}_{4}$ in water are $4.7 \times 10^{-9}$ and $1.3 \times 10^{-9}$, respectively, at $25^{\circ} \mathrm{C}$. If a mixture of two is washed with $\mathrm{H}_{2} \mathrm{O}$, what is $\mathrm{Ca}^{2+}$ ion concentration in water?
A. $7.7 \times 10^{-5}$
B. $5.831 \times 10^{-5}$
C. $6.856 \times 10^{-5}$
D. $3.606 \times 10^{-5}$

## Answer: A

## - Watch Video Solution

84. What are the units in which the solubility product of $\mathrm{Ca}_{3}\left(\mathrm{PO}_{4}\right)_{2}$ is expressed?
A. moldm $^{-3}$
B. $\mathrm{mol}^{2} \mathrm{dm}^{-6}$
C. $\mathrm{mol}^{3} \mathrm{dm}^{-9}$
D. $\mathrm{mol}^{5} \mathrm{dm}^{-15}$

## Answer: D

85. The Ph of $10^{-5} \mathrm{MHCl}$ solution if 1 " mL of " it is diluted to 1000 mL is
A. 5
B. 8
C. 7.02
D. 6.98

## Answer: D

## - Watch Video Solution

86. Which of the following when mixed, will given a solution with $\mathrm{pH}>7$.
A. $0.1 \mathrm{MHCl}+0.1 \mathrm{MNaOH}$
B. $100 \mathrm{mLof0} .1 \mathrm{MH}_{2} \mathrm{SO}_{4}+100 \mathrm{mLof0} .3 \mathrm{MNaOH}$
C. $100 \mathrm{mLof0} 0.1 \mathrm{MHC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+100 \mathrm{mLof0} 0.1 \mathrm{MKOH}$
D. $25 \mathrm{mLof0} .1 \mathrm{HNO}_{3}+25 \mathrm{mLof0} .1 \mathrm{MNH}_{3}$

## Answer: C

## - Watch Video Solution

87. A solution of $\mathrm{CaF}_{2}$ is found to contain $4 \times 10^{-4} \mathrm{M}$ of $F^{\Theta}, K_{s p}$ of $\mathrm{CaF}_{2}$ is
A. $3.2 \times 10^{-11}$
B. $0.8 \times 10^{-11}$
C. $6.4 \times 10^{-11}$
D. $32 \times 10^{-11}$

## Answer: A

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88. At what $p H$ will a $1.0 \times 10^{-3} \mathrm{M}$ solution of an indicator with $K_{b}=1.0 \times 10^{-10}$ changes colour?
A. 10
B. 4.0
C. 3
D. 7

## Answer: B

## - Watch Video Solution

89. $K_{b}$ for $\mathrm{NH}_{4} \mathrm{OH}$ is $1.8 \times 10^{-5}$. The $\left[\begin{array}{l}\Theta \\ \mathrm{OH}\end{array}\right]$ of $0.1 \mathrm{MNH}_{4} \mathrm{OH}$ is
A. $1.8 \times 10^{-6}$
B. $1.34 \times 10^{-3}$
C. $4.20 \times 10^{-2}$
D. $5.0 \times 10^{-2}$

## Answer: B

90. $p H$ signifies:
A. Puissance de hydrogen
B. $\log \left[H^{+}\right]$
C. $-14-\mathrm{pOH}$
D. All the above

## Answer: A

## - Watch Video Solution

91. A solution with $p H=2$ is more acidic than one with a $p H=6$ by a factor of
A. 4
B. 12
C. 400
D. $10^{4}$

## Answer: D

## - Watch Video Solution

92. A definite volume of an aqueous $N / 20$ acetic acid $\left(p K_{a}=4.74\right)$ is titrated with a strongs base. It is found that 75 equal-sized drops of NaOH added from a burette effect the complete neutralisation. Find the $p H$ when an acid solution is neutralised to the extent of $20 \%, 40 \%$, and $90 \%$, respectively.
A. 4.14
B. 9.86
C. 5.34
D. 8.68

## D Watch Video Solution

93. The $p K_{a}$ of acteylsalicylic acid (aspirin) is 3.5 . The pH of gastric juice in human stomach is about 2-3 and the pH in the small intestine is about 8. Aspirin will be:
A. Unionised in the small intestine and in the stomach.
B. Completely ionised in the small intestine and in the stomach.
C. Ionised in the stomach and alomost unionised in the small intestine.
D. lonised in small intestine and almost unionised in the stomach.

## Answer: D

## D Watch Video Solution

94. Which of the following salt is basic?
A. HOCI
B. NaOCI
C. $\mathrm{NaHSO}_{4}$
D. $\mathrm{NH}_{4} \mathrm{NO}_{3}$

## Answer: B

## - Watch Video Solution

95. For the indicator 'HIn' the ratio $\left(\right.$ Ind $\left.^{\Theta}\right) /(H I n)$ is 7.0 at $p H$ of 4.3. What is $K_{e q}$ for the indicator.
A. $3.5 \times 10^{-4}$
B. $3.5 \times 10^{-5}$
C. $3.5 \times 10^{-2}$
D. $3.5 \times 10^{-3}$

## Answer: A

## - Watch Video Solution

96. When 0.002 mol of acid is added to 250 mL of a buffer solution, pH decreases by 0.02 units. The buffer capacity of the system is
A. (a) 0.1
B. (b) 0.2
C. (c) 0.3
D. (d) 0.4

## Answer: D

## - Watch Video Solution

97. pH of an aqueous solution of $0.6 \mathrm{MNH}_{3}$ and $0.4 \mathrm{MNH}_{4} \mathrm{Cl}$ is $9.4\left(p K_{b}=4.74\right)$. The new $p H$ when $0.1 M \mathrm{Ca}(\mathrm{OH})_{2}$ solution is added to it.
A. 9.86
B. 10.14
C. 10.2
D. 10.86

## Answer: A

## - Watch Video Solution

98. Which of the following salts undergoes anionic hydrolysis?
A. (a) $\mathrm{CuSO}_{4}$
B. (b) $\mathrm{NH}_{4} \mathrm{Cl}$
C. (c) $\mathrm{FeCl}_{3}$
D. (d) $\mathrm{Na}_{2} \mathrm{CO}_{3}$

## Answer: D

## - Watch Video Solution

99. A saturated solution of $\mathrm{Ag}_{2} \mathrm{SO}_{4} i s 2.5 \times 10^{-2} M$. The value of its solubility product is
A. $62.5 \times 10^{-6}$
B. $6.25 \times 10^{-4}$
C. $15.625 \times 10^{-10}$
D. $3.125 \times 10^{-6}$

## Answer: A

## - Watch Video Solution

100. Which one of the followinf is acid salt?
A. $\mathrm{Na}_{2} \mathrm{~S}$
B. $\mathrm{Na}_{2} \mathrm{SO}_{3}$
C. $\mathrm{NaHSO}_{3}$
D. $\mathrm{Na}_{2} \mathrm{SO}_{4}$

## Answer: C

## - Watch Video Solution

101. Which one is not an acid salt?
A. $\mathrm{NaH}_{2} \mathrm{PO}_{4}$
B. $\mathrm{NaH}_{2} \mathrm{PO}_{2}$
C. $\mathrm{NaH}_{2} \mathrm{PO}_{3}$
D. All of the above are acid salts

## Answer: D

## D Watch Video Solution

102. Which one of the following salts when dissolves in water hydrolyse?
A. NaCI
B. $\mathrm{NH}_{4} \mathrm{CI}$
C. $K C I$
D. $\mathrm{Na}_{2} \mathrm{SO}_{4}$

## Answer: B

103. Which of the following salt undergoes hydrolysis?
B. $\mathrm{NaNO}_{3}$
C. KCI
D. $\mathrm{K}_{2} \mathrm{SO}_{4}$

## Answer: A

## - Watch Video Solution

104. Out of following the compound whose water solution has the highest $p H$ is
A. NaCI
B. $\mathrm{Na}_{2} \mathrm{CO}_{3}$
C. $\mathrm{NH}_{4} \mathrm{CI}$
D. $\mathrm{NaHCO}_{3}$

## Answer: B

105. When equal volumes of the following solutions are mixed, the precipitation of $\mathrm{AgCl}\left(K_{s p}=1.8 \times 10^{-10}\right)$ will occur with :
A. $10^{-4} \mathrm{M}\left(\mathrm{Ag}{ }^{\oplus}\right)$ and $10^{-4} \mathrm{M}\left(\mathrm{CI}^{\Theta}\right)$
B. $10^{-5} \mathrm{M}\left(\mathrm{Ag}^{\oplus}\right)$ and $10^{-5} \mathrm{M}\left(\mathrm{CI}^{\Theta}\right)$
C. $10^{-5} \mathrm{M}\left(\mathrm{Ag}^{\oplus}\right)$ and $10^{-6} \mathrm{M}\left(\mathrm{CI}^{\Theta}\right)$
D. $10^{-4} \mathrm{M}\left(A g^{\oplus}\right)$ and $10^{-10} \mathrm{M}\left(C I^{\Theta}\right)$

## Answer: A

## - Watch Video Solution

106. The gastric juice in our stomach contains enough HCl to make the hydrogen ion concentration about $0.01 \mathrm{~mol} \mathrm{~L}^{-1}$. The pH of gastric juice is
A. (a) 0.01
B. (b) 1
C. (c) 2
D. (d) 14

## Answer: C

## - Watch Video Solution

107. Of the given anions, the strongest Bronsted base is
A. (a) $\mathrm{ClO}^{\Theta}$
B. (b) $\mathrm{ClO}_{2}^{\Theta}$
C. (c) $\mathrm{ClO}_{3}^{\Theta}$
D. (d) $\mathrm{ClO}_{4}^{\Theta}$

## Answer: A

108. In decinormal solution, $\mathrm{CH}_{3} \mathrm{COOH}$ acid is ionised to the extent of $1.3 \%$. If $\log 1.3=0.11$, what is the pH of the solution?
A. 3.89
B. 2.89
C. 4.89
D. Unpredictable

## Answer: B

## - Watch Video Solution

109. An aqueous solution of aluminium sulphate would show
A. Acidic
B. Neutral
C. Basic
D. Both acidic and basic reaction.

## D Watch Video Solution

110. AICI $_{3}$ achieves stability by forming a dimer. In trivalent state the compound is hydrolysed in water. $\mathrm{AICI}_{3}$ in acidified aqueous solution forms
A. Cation hydrolysis
B. Anion hydrolysis
C. Hydrolysis of both anion and cation
D. Dissociation

## Answer: A

111. A solution contains 10 mL of 0.1 NNaOH and 10 mL of $0.05 \mathrm{Na}_{2} \mathrm{SO}_{4}, \mathrm{pH}$ of this solution is
A. 7
B. Less than 7
C. Greater than 7
D. Zero

## Answer: C

## - Watch Video Solution

112. 20 mL of 0.1 NHCI is mixed with 20 ml of 0.1 NKOH . The pH of the solution would be
A. 0
B. 7
C. 2
D. 9

## Answer: B

## - Watch Video Solution

113. 0.1 M solution of which one of these substances will be basic ?
A. Sodium borate
B. Ammonium ditoride
C. Calcium nitrate
D. Sodium sulphate

## Answer: A

## - Watch Video Solution

114. In which of the following solvents will AgBr has highest solubility?
A. (a) $10^{-3} \mathrm{MNaBr}$
B. (b) $10^{-3} \mathrm{MNH}_{4} \mathrm{OH}$
C. (c) Pure water
D. (d) $10^{-3} \mathrm{MHBr}$

## Answer: B

## - Watch Video Solution

115. Which of the following solutions will have pH close to 1.0 ?
A. $100 \mathrm{mLM} / 10 \mathrm{HCI}+100 \mathrm{mLM} / 10 \mathrm{NaOH}$
B. $55 \mathrm{mLM} / 10 \mathrm{HCI}+45 \mathrm{mLM} / 10 \mathrm{NaOH}$
C. $10 \mathrm{mLM} / 10 \mathrm{HCI}+90 \mathrm{mLM} / 10 \mathrm{NaOH}$
D. $75 \mathrm{mLM} / 5 \mathrm{HCI}+25 \mathrm{mLM} / 5 \mathrm{NaOH}$

## Answer: D

116. Fear or excitement, generally cause one to breathe rapidly and it results in the decrease of concentration of $\mathrm{CO}_{2}$ in blood. In what way it will change pH of blood ?
A. $p H$ will increase
B. $p H$ will decrease
C. No change
D. pH will adjust to 7

## Answer: C

## - Watch Video Solution

117. Which buffer solution out of the following will have $\mathrm{pH}>7$ ?
A. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{CH}_{3} \mathrm{COONa}$
B. $\mathrm{HCOOH}+\mathrm{HCOOK}$
C. $\mathrm{CH}_{3} \mathrm{COONH}_{4}$
D. $\mathrm{NH}_{4} \mathrm{OH}+\mathrm{NH}_{4} \mathrm{CI}$

## Answer: D

## - Watch Video Solution

118. Which of the following is most soluble ?
A. $B i_{2} S_{3}\left(K_{s p}=1 \times 10^{-70}\right)$
B. $\operatorname{MnS}\left(K_{s p}=7 \times 10^{-16}\right)$
C. $\operatorname{CuS}\left(K_{s p}=8 \times 10^{-37}\right)$
D. $A g_{2} S\left(K_{s p}=6 \times 10^{-51}\right)$

## Answer: B

## - Watch Video Solution

119. If $\mathrm{H}_{3} \mathrm{O}^{+}$ion concentration of a solution is increased by 10 times, its pH will
A. Increase by 1
B. Remains unchanged
C. Decreases by 1
D. Increase by 10

## Answer: C

## - Watch Video Solution

120. If $p K_{b}$ for fluoride ion at $25^{\circ} \mathrm{C}$ is 10.83 the ionisation constant of hydrofluoric acid in water at this temperature is
A. $1.74 \times 10^{-5}$
B. $3.52 \times 10^{-3}$
C. $6.75 \times 10^{-4}$
D. $5.38 \times 10^{-2}$

Answer: C

## - Watch Video Solution

121. The following graph represents the titration of pH vs volume

A. A diprotic acid.
B. Two monoprotic acids with the same $K_{a}$ but different concentrations.
C. Two monoprotic acids with different $K_{a}$ but the same concentration.
D. Two monoprotic acids with differnet $K_{a}$ and different concentartions.

## Answer: D

## - Watch Video Solution

## Exercises Assertion-Reasoning

1. Assertion (A): A solution contains 0.1 M each of $\mathrm{pB}^{2+}, \mathrm{Zn}^{2+}, \mathrm{Ni}^{2+}$, ions.

If $\mathrm{H}_{2} \mathrm{~S}$ is passed into this solution at $25^{\circ} \mathrm{C}$.
$\mathrm{Pb}^{2+}, \mathrm{Ni}^{2+}, \mathrm{Zn}^{2+}$ will get precpitated simultanously.
Reason (R): $\mathrm{Pb}^{2+}$ and $\mathrm{Zn}^{2+}$ will get precipitated if the solution contains

### 0.1MHCI.

$$
\left[K_{1} H_{2} S=10^{-7}, K_{2} H_{2} S=10^{-14}, K_{s p} P b S=3 \times 10^{-29} K_{s p} N i S=3 \times 10^{-19} \cdot K_{s p} Z n!\right.
$$

A. If both (A) and (R) are correc, and (R) is the correct explanation of

## (A).

B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## - Watch Video Solution

2. Assertion (A): Solubility of $A g C N$ in acidic solutions is greater than in pure water.

Reason (R) : Solubility equilibrium of $A g C N$ is shifted in forward direction due to the formation of $H C N$.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but ( $R$ ) is correct.

## Answer: A

## - Watch Video Solution

3. Methy1 red has a $K_{a}=10^{-5}$. The acid form Hin is red and its conjugate base, $I n d^{\Theta}$ is yellow. Complete the following table:
$\begin{array}{llll}\mathrm{pH} & 3 & 5 & 7\end{array}$
$\left[\right.$ Ind $\left.^{\Theta}\right] /[\mathrm{HIn}] \quad-\quad-$
Colour
A. If both (A) and (R) are correc, and (R) is the correct explanation of

## (A).

B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## D Watch Video Solution

4. Assertion (A) : On cooling in a freezing mixture, colout of the mixture turns to pink from deep blue for a reaction.
$\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}^{2+}($ Pink $)(\mathrm{aq})+4 \mathrm{CI}^{\Theta} \Leftrightarrow \mathrm{CoCI}_{4}^{2-}(a q)($ Blue $)+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
Reason (R) : The reaction is endothermic in forward reaction,so on cooling the reaction, deep blue colour appears.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## D Watch Video Solution

5. Assertion (A): Due to common ion effect, the solubility of $\mathrm{HgI}_{2}$ is expected to be less in an aqueous solution of KI than in water. But $\mathrm{HgI}_{2}$ dissolves in an aqueous solution of $K I$ to form a clear solution. Reason (R) : $I^{\Theta}$ ions is highly polarisable.
A. If both (A) and (R) are correc, and (R) is the correct explanation of
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: B

## - Watch Video Solution

6. Assertion (A): $p K_{a}$ of a weak acid become equal of the $p H$ of the solution at the mid-point of titration.

Reason (R) : The molar concentration of the proton donor an proton acceptor beomes equal at the mid-point.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: B

## - Watch Video Solution

7. Assertion (A): pH of HCI solution is less than that of acetic acid of the some concentartion.

Reason (R) : In equimolar solution, the number of titrable protons present in $H C I$ is less than that present in acetic acid.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

## - Watch Video Solution

8. Assertion (A): $p H$ value of $H C N$ solution decreases when $N a C N$ is added to it.

Reason (R) : NaCN provides a common ion $C N^{\Theta}$.
A. If both (A) and (R) are correc, and (R) is the correct explanation of
(A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: D

## - Watch Video Solution

9. Assertion (A): pH of water increases with an increase in temperature.

Reason (R) : $K_{w}$ or water increases with increase in temperature.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: D

## - Watch Video Solution

10. Assertion (A): $\mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}$ is a salt of unstable acid.

Reason (R) : $\mathrm{H}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}$ is a polyprotic acid.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: B

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11. Assertion (A): When a solution of $\mathrm{CH}_{3} \mathrm{COOH}$ in water is shaken with charcoal, pH of the solution will get decreased.

Reason $(\mathrm{R})$ : The degree of ionisation of $\mathrm{CH}_{3} \mathrm{COOH}$ increase.
A. If both (A) and (R) are correc, and (R) is the correct explanation of
(A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: D

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12. Assertion (A): There is very little difference in acid strength of $\mathrm{H}_{3} \mathrm{PO}_{4}, \mathrm{H}_{3} \mathrm{PO}_{3}$, and $\mathrm{H}_{3} \mathrm{PO}_{2}$.

Reason (R) : The hydrogens in these acids are not all bonded to oxygens. The electrone-grativities of $P$ and $H$ are almost the same.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## - Watch Video Solution

13. Assertion (A): Both reactions are Lewis acid-base recations?
i. $\mathrm{NH}_{3}+\mathrm{BF}_{3} \rightarrow \mathrm{H}_{3} \mathrm{~N}: \mathrm{BF}_{3}$
ii. $\mathrm{Mg}+\mathrm{S} \rightarrow \mathrm{Mg}^{2+}+\mathrm{S}^{2-}$

Reason (R) : Lewis acid-base reaction involve the donation of lone pair electrons from base to acid. this donation results in a corrdinate bond.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both $(A)$ and $(R)$ are correct but $(R)$ is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

Answer: D

## - Watch Video Solution

14. Assertion (A): Solution of $\mathrm{AICI}_{3}$ in water is neutral.

Reason (R) : $\left[A I\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}$ is formed.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: D

15. Assertion (A): In dilute benzene solutions, equimolar addition of $R_{3} N$ and HCI produce a substance with a dipole moment. In the same solvent, equimolar addition of $R_{3} \mathrm{~N}$ and $\mathrm{SO}_{3}$ produce a substance having an almost identical dipole moment.

Reason (R) : Both HCI and $\mathrm{SO}_{3}$ are Lewis acids and can react with the amine base to form polar substances which undergo ionic dissociation in a solvent sufficiently more polar than benzene.

Moreover, $(N-S)$ bond is a more polar bond.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## (D) Watch Video Solution

16. Assertion (A): A certain recation is catalysed by acids and the catalytic activity of 0.1 M solutions of the acids in water decrease in the order, $\mathrm{HCI}, \mathrm{HCOOH}$, and $\mathrm{CH}_{3} \mathrm{COOH}$. The same reaction takes place in anhydrous $\mathrm{NH}_{3}$, but the three acids have same catalytic effect in 0.1 M solution.

Reason (R) : The order of catalytic activity in water is the same as the order of acidity. in anhyrous $\mathrm{NH}_{3}$, all the three acids are strong.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both $(A)$ and $(R)$ are correct but $(R)$ is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## Watch Video Solution

17. Assertion (A): S reacts with $\mathrm{SO}_{3}^{2-}$ and forms $\mathrm{S}_{2} \mathrm{O}_{3}^{2-}$

Reason (R) : $S$ is electorn deficient and acts and acid and $\mathrm{SO}_{3}^{2-}$ is a base in terms of Lewis acid theroy.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## - Watch Video Solution

18. Assertion (A): The amino acid glycine predominatly exists in the form of $\wedge(\oplus) \mathrm{NH}_{3} \mathrm{CH}_{2} \mathrm{COO}^{\Theta}$.

Reason (R) : The conjugate acid of glycine is $\mathrm{NH}_{2} \mathrm{CH}_{2} \mathrm{COO}^{\Theta}$.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but ( $R$ ) is correct.

## Answer: C

## - Watch Video Solution

19. Assertion (A): $\mathrm{Sb}^{3+}$ is not precipitated as sulphide when $\mathrm{H}_{2} \mathrm{~S}(\mathrm{~g})$ is passed in alkaline solution.

Reason $(R):\left[S^{2-}\right]$ ion in basic medium is inadequate for precipitation.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

## - Watch Video Solution

20. Assertion (A): Addition of $\mathrm{Ag}^{\oplus}$ ions to a mixture of aqueous NaCI and

NaBr solution will first precipitate AgBr rather than AgCI .
Reason (R): $K_{s p} A g C I<K_{s p} o f A g B r$.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

## - Watch Video Solution

21. Assertion : The pH of an aqueous solution of acetic acid remains unchanged on addition of sodium acetate.

Reason : The ionization of acetic acid is increased by addition of sodium acetate.
A. If both (A) and (R) are correc, and (R) is the correct explanation of
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: D

## - Watch Video Solution

22. Assertion (A): On mixing equal volumes of 1 MHCi and $2 \mathrm{MCH}_{3} \mathrm{COONa}$, an acidic buffer solution is formed.

Reason (R) : The resultant mixture contains $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COONa}$ which are parts of acidic buffer.
A. If both (A) and (R) are correc, and (R) is the correct explanation of

> (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

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23. Assertion (A): On addition of $\mathrm{NH}_{4} \mathrm{CI}$ to $\mathrm{NH}_{4} \mathrm{OH}, \mathrm{pH}$ decreases but remains greater than 7 .

Reason (R) : Addition of $\stackrel{\oplus}{\mathrm{NH}_{4}}$ ion decreases ionisation of $\mathrm{NH}_{4} \mathrm{OH}$, thus
$\left[\begin{array}{l}\mathrm{O} \\ \mathrm{OH}\end{array}\right]$ decreases and also pH decreases.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of
(A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

## - Watch Video Solution

24. Assertion (A): A is very dilute acidic solution of $\mathrm{Cd}^{2+}$ and $\mathrm{Ni}^{2+}$ gives yellow precipitate of CdS on passing hydrogen sulphide. Reason (R) : Solubility product of CdS is more than that of NiS.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but ( $R$ ) is correct.

## Answer: C

25. STATEMENT-1: In the titratio of $\mathrm{Na}_{2} \mathrm{CO}_{3}$ with HCl using methyl orange indicator, the volume of the acid required at the equvalence point is twice that of the acid required using phenolphthalenin as inkdicator.

STATEMENT-2: Two moles of HCl are required for the complete neutralisation of one mole of $\mathrm{Na}_{2} \mathrm{CO}_{3}$
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: B

26. Statement: In acidic medium, $\mathrm{Zn}^{2+}$ is not precipitated by $\mathrm{H}_{2} \mathrm{~S}$.

Explanation: Common ion effect reduces the concentration of $S^{2-}$ to a minimum level.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

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27. Statement: In an acid-basic titration involving a strong base and a weak acid, methyl orange can be used as an indicator.

Explanation: Methyl orange changes its colour in the pH range 3 to 5.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If both (A) and (R) are incorrect.

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28. Assertion (A): An aqueous solution of ammonium acetate acts as a buffer solution.

Reason (R) : A buffer solution reacts with small quantities of hydrogen or hydroxy ions and keps the pH almost same.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: B

## D Watch Video Solution

29. Assertion (A): When small amount of acid or base is added to pure water, its pH undergoes a change.

Reason (R) : Addition of an acid or a basic increases the degree of ionisation of water.
A. If both (A) and (R) are correc, and (R) is the correct explanation of
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

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30. Assertion (A): pH of acidic solution is always below 7 at $25^{\circ} \mathrm{C}$. Reason ( R ) : At $25^{\circ} \mathrm{C}$, the pH of $10^{-8} \mathrm{MHCI}$ is 8 .
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

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31. Assertion (A): $\mathrm{H}_{2} \mathrm{SO}_{4}$ acts as a base in the presence of $\mathrm{HCIO}_{4}$.

Reason (R) : Perchloride acid is stronger acid than $\mathrm{H}_{2} \mathrm{SO}_{4}$.
A. If both $(A)$ and $(R)$ are correc, and $(R)$ is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: A

32. Assertion (A): $p H$ of neutral solution is always 7 .

Reason ( R ) : pH of solution does not depend upon temperature.
A. If both $(A)$ and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If both (A) and (R) are incorrect.

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33. Assertion (A): pH of $10^{8} \mathrm{MHCl}$ is not equal to 8 .

Reason (R) : HCl does not dissociate properly in very dilute solution.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

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34. Assertion (A): If a solution with $p H=2$ is diluted to double the volume, the pH of the solution will fall to 1 .

Reason ( R ) : $p H$ is inversely proportional to the volume of the solution.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If both $(A)$ and $(R)$ are incorrect.

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35. Assertions : When HCl gas is passed through a saturated solution of common salt, pure NaCl is precipitated.

Reason: The solubility product of NaCl gets lowered.
A. If both (A) and (R) are correc, and (R) is the correct explanation of (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but (R) is correct.

## Answer: C

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36. Assertion (A): pH of buffer chnages with temperature.

Reason (R) : Ionic of a water $\left(K_{w}\right)$ changes with temperature.
A. If both (A) and (R) are correc, and (R) is the correct explanation of

> (A).
B. If both (A) and (R) are correct but (R) is not the correct explanation of (A).
C. If (A) is correct, but (R) is incorrect.
D. If (A) is incorrect, but ( $R$ ) is correct.

## Answer: A

## Exercises Integer

1. What is the sum of magic numbers of all solutions gives below : (Interger value is between 50 and 60)
(Magic numver of a solution $=p H$ of solution $\times$ Weight factor)

Solution
I $\quad 0.1 \mathrm{M} \operatorname{HCN}\left(K_{a}=10^{-10}\right)$
II $0.1 \mathrm{MCH}_{3} \mathrm{COOH}+0.1 \mathrm{MCH}_{3} \mathrm{COONa}\left(\mathrm{K}_{a}=10^{-5}\right)$
III 0.1 MHCl
IV $0.1 \mathrm{MNH}_{4} \mathrm{OH}\left(K_{b}=10^{-5}\right)$
V 0.01 MNaOH 0.5

VI 10 mL of0.01 $\mathrm{MCH}_{3} \mathrm{COOH}+10 \mathrm{ml}$ of $0.1 \mathrm{MNH}_{4} \mathrm{OH} \quad 1$
2
1
3

2

Weight factor

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(-) Watch Video Solution
2. How much of the following cations belong to group IIA, III, IV, and V only in qualitative salt analysis?
$\mathrm{Sr}^{2+}, \mathrm{Hg}_{2}^{2+}, \mathrm{Pb}^{2+}, \mathrm{Zn}^{2+}, \mathrm{Cu}^{2+}, \mathrm{Cr}^{3+}, \mathrm{Ag}^{+}, \mathrm{Mg}^{2+}, \mathrm{Sb}^{3+}$
3. How much of the following are strong electrolytes?
a. $\mathrm{NH}_{3}$ b. $\mathrm{NH}_{4} \mathrm{Cl}$
c. $\mathrm{CH}_{3} \mathrm{COOH}$
d. $\mathrm{CH}_{3} \mathrm{COONa}$ e. HCl f. NaCl

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4. How much of the following 0.1 M solutions are acidic?
a. $\mathrm{NH}_{4} \mathrm{CI}$ b. NaOH c. $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ d. NaCI
e. $\mathrm{NH}_{3}+\mathrm{NH}_{4} \mathrm{CI}$ f. $\mathrm{NH}_{3}$ g. HCI
h. $\mathrm{HCIO}_{4}$ i. $\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4}$ j. $\mathrm{K}_{2} \mathrm{SO}_{4}$

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5. How many are basic ?
6. How many compounds are neutral ? $\mathrm{H}_{2} \mathrm{O}, \mathrm{NaOH}, \mathrm{HCL}, \mathrm{Nacl}$

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7. How many of the following salts:
i. $\mathrm{NH}_{4} \mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ ii. $\mathrm{PhCOONH}_{4}$ iii. $\mathrm{NaC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$
iv. $\mathrm{NH}_{4} \mathrm{CI}$ v. MgS vi. $\mathrm{Na}_{2} \mathrm{SO}_{4}$
vii. KCI
a. Hydrolyse more in water at $25^{\circ} \mathrm{C}$.
b. Do not hydrolyse.

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8. How many of the following combinations of reactants will react less than $2 \%$ of theroetically possible extent?
a. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{H}_{2} \mathrm{O}$ b. $\mathrm{CH}_{3} \mathrm{COO}^{\Theta}+\mathrm{H}_{2} \mathrm{O}$
c. $\mathrm{CH}_{3} \mathrm{COO}^{\Theta}+\mathrm{H}_{3} \mathrm{O}^{\oplus}$ d. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{KOH}$

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9. How many of the conbinations of reactants in above will react untill more than $98 \%$ of the limiting quantity is used up?
a. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{H}_{2} \mathrm{O}$ b. $\mathrm{CH}_{3} \mathrm{COO}^{\Theta}+\mathrm{H}_{2} \mathrm{O}$

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10. Calculate the $p H$ at equilibrium point when a solution of $10^{-6} \mathrm{MCH}_{3} \mathrm{COOH}$ is titrated with a solution of $10^{-6} \mathrm{MNaOH} . \mathrm{K}_{a}$ for acid $2 \times 10^{-5}\left(p K_{a}=4.7\right)$ (Answer given in whole number).

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## Exercises True/ False

1. Silver chloride is more soluble in very concentrated sodium chloride solution than in pure water. (T/F)
2. Any buffer solution can be used as a buffer upto two pH units only.

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3. $\mathrm{Mg}^{2+}$ ions is essential for selective precipitation of $\mathrm{Fe}(\mathrm{OH})_{3}$ by aqueous $\mathrm{NH}_{3}$.

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4. A mixture of aqueous solution of sodium acetate and sodium propanoate forms a buffer solution. (T/F)

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5. pH of some solution is given by $p H=\frac{p K_{a_{1}}+p K_{a_{2}}}{2}$. This formula is valid for the compound $\mathrm{NaH}_{2} \mathrm{BO}_{3}$.

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6. 0.6 mmol of NaCl and 1 mol of HCI in 1 L solution is a buffer.

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7. The $\mathrm{K}_{a}$ for $\mathrm{CH}_{3} \mathrm{COOH}$ at 300 and 310 K are $1.8 \times 10^{-5}$ and $1.805 \times 10^{-5}$, respectively. The enthalpy of deprotonation for acetic acid is 51.6cal.

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8. Out of the following salts:
i. $\mathrm{NaH}_{2} \mathrm{BO}_{3}$ ii. $\mathrm{NH}_{2} \mathrm{CH}_{2} \mathrm{COOH}$
iii. $\mathrm{CH}_{3} \mathrm{COONH}_{4}$ iv. NaHS

Salts (i) has concentration-depends pH .

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9. Out of the following acid base-reactions, reaction
(b) and (c) are possible.
a. $\mathrm{PH}_{3}+\stackrel{\oplus}{\mathrm{N}} \mathrm{H}_{4} \rightarrow \stackrel{\oplus}{\mathrm{P}} \mathrm{H}_{4}+\mathrm{NH}_{3}$
b. $\mathrm{NH}_{3}+\stackrel{\oplus}{\mathrm{P}}{ }_{4} \rightarrow \stackrel{\oplus}{\mathrm{NH}} \mathrm{H}_{4}+\mathrm{PH}_{3}$
c. $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{P}+\stackrel{\oplus}{\mathrm{N}} \mathrm{H}_{4} \rightarrow\left(\mathrm{CH}_{3}\right)_{3} \stackrel{\oplus}{\mathrm{P}} \mathrm{H}+\mathrm{NH}_{3}$
d. $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{~N}+\mathrm{PH}_{4}^{\oplus} \rightarrow\left(\mathrm{CH}_{3}\right)_{2}{ }^{\oplus} \mathrm{NH}+\mathrm{PH}_{3}$

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10. The oxo-acids of $P_{2} \mathrm{O}_{5}$ is $\mathrm{H}_{3} \mathrm{PO}_{4}$

## - Watch Video Solution

1. Which of the following statements is//are correct?
A. The $p H$ of $1.0 \times 10^{-8} \mathrm{M}$ solution of $H C I$ is 8 .
B. The conjugate base of $\mathrm{H}_{2} \mathrm{PO}_{4}^{\Theta}$ is $\mathrm{HPO}_{4}^{-2}$.
C. The autoprotolysis constant of water increases with temperature.
D. When a solution of a weak monoprotic acid is trated against a strong base, at half-neutralisation point, $p H=(1 / 2) p K_{a}$.

## Answer: B::C

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2. A buffer solution can be prepared from a mixture of
A. Sodium acetate and acetic acid in water.
B. Sodium acetat and HCI in water
C. Ammonia and ammonia chloride in water.
D. Ammonia and sodium hydroxide in water.

## Answer: A::C

## D Watch Video Solution

3. Aqueous solution of $\mathrm{HNO}_{3}, \mathrm{CH}_{3}, \mathrm{CH}_{3} \mathrm{COOH}$, and $\mathrm{CH}_{3} \mathrm{COOK}$ of identical concentrations are given. The pair (s) of the solution which may form a buffer upon mixing is (are):
A. $\mathrm{HNO}_{3}$ and $\mathrm{CH}_{3} \mathrm{COOH}$
B. KOH and $\mathrm{CH}_{3} \mathrm{COONa}$
C. $\mathrm{HNO}_{3}$ and $\mathrm{CH}_{3} \mathrm{COONa}$
D. $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COONa}$

## Answer: C::D

## Archives Single Correct

1. At $90^{\circ} \mathrm{C}$, pure water has $\left[\mathrm{H}_{3} \mathrm{O}^{\oplus}\right]=10^{-7} \mathrm{molL}{ }^{-1}$. What is the value of $K_{w}$ at $90^{\circ} \mathrm{C}$ ?
A. $10^{-6}$
B. $10^{-12}$
C. $10^{-14}$
D. $10^{-8}$

## Answer: B

## - Watch Video Solution

2. Calculate pH of $1.0 \times 10^{-8} \mathrm{M}$ solution of HCl .
A. 8
B. -8
C. Between 7 and 8
D. Between 6 and 7

## Answer: D

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3. An acidic buffer solution can be prepared by mixing the solution of
A. Sodium acetate and acetic acid
B. Ammonium chloride and ammonium hydroxide
C. Sulpuric acid and sodium sulphate
D. Sodium chloride and sodium hydroxide

## Answer: A

4. The strongest Bronsted base in the following anion is
A. $\mathrm{CIO}^{\Theta}$
B. $\mathrm{CIO}_{2}{ }^{\Theta}$
C. $\mathrm{CIO}_{3}{ }^{\Theta}$
D. $\mathrm{CIO}_{4}^{\Theta}$

## Answer: A

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5. The precipitate of $\mathrm{CaF}_{2}$ is obtained when equal volumes of the following are mixed.
$\left[K_{s p}\left(\mathrm{CaF}_{2}\right)=1.7 \times 10^{-10}\right]$
A. $10^{-4} \mathrm{MCa}^{2+}+10^{-4} \mathrm{MF}^{\Theta}$
B. $10^{-2} \mathrm{MCa}^{2+}+10^{-3} \mathrm{MF}^{\Theta}$
C. $10^{-5} \mathrm{MCa}^{2+}+10^{-3} \mathrm{MF}^{\Theta}$
D. $10^{-3} \mathrm{MCa}^{2+}+10^{-5} \mathrm{MF}^{\Theta}$

## Answer: B

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6. A certain buffer solution contains equal concentartion of $X^{\Theta}$ and $H X$. The $K_{b}$ for $X^{\Theta}$ is $10^{-10}$. The $p H$ of the buffer is
A. 4
B. 7
C. 10
D. 14

## Answer: A

7. $K_{a}$ of $H A$ is $10^{-4}$. The equilibrium constant for its reaction with a strong base is
A. $10 \times 10^{-4}$
B. $10 \times 10^{-10}$
C. $10 \times 10^{10}$
D. $1.0 \times 10^{14}$

## Answer: C

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8. Conjugate acid of $\mathrm{NH}_{2}$ :
A. $\mathrm{NH}_{3}$
B. $\mathrm{NH}_{2} \mathrm{OH}$
C. $\mathrm{NH}_{4}^{\oplus}$
D. $\mathrm{N}_{2} \mathrm{H}_{4}$

## D Watch Video Solution

9. The best indicator for the detection of end point in titration of a weak acid and a strong base is :
A. Methy1 orange $(3 \rightarrow 4)$
B. Methy1 red $(5 \rightarrow 6)$
C. Bromothymol blue ( $6 \rightarrow 7.5$ )
D. Phenolphtalein (8 $\rightarrow$ 9.6)

## Answer: D

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10. The compound that is not a Lewis acids is
A. $B F_{3}$
B. $\mathrm{AICI}_{3}$
C. $\mathrm{BeCI}_{2}$
D. $\mathrm{SnCI}_{4}$

## Answer: D

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11. The $p K_{a}$ of acety1 salicylic acid (aspirin) is 3.5. The $p H$ of gastric juice in human stomach is about 2.3 and the $p H$ in the same intestine is bout 8 . A spirin will be
A. Unionised in the small intestine and in the stomach.
B. Completely ionised in the small intestine and in the stomach.
C. lonised in the stomach and alomost unionised in the small intestine.
D. lonised in the small intestine and almost unionised in the stomach.

## Answer: D

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12. When equal volumes of the following solutions are mixed, the precipitation of $\mathrm{AgCl}\left(K_{s p}=1.8 \times 10^{-10}\right)$ will occur with :
A. $10^{-4} \mathrm{M}\left(\mathrm{Ag}^{\oplus}\right)$ and $10^{-4} \mathrm{M}\left(\mathrm{CI}^{\Theta}\right)$
B. $10^{-5} \mathrm{M}\left(\mathrm{Ag}^{\oplus}\right)$ and $10^{-5} \mathrm{M}\left(\mathrm{CI}^{\Theta}\right)$
C. $10^{-6} \mathrm{M}\left(\mathrm{Ag}^{\oplus}\right)$ and $10^{-6} \mathrm{M}\left(C I^{\Theta}\right)$
D. $10^{-10} \mathrm{M}\left(A g^{\oplus}\right)$ and $10^{-10} \mathrm{M}\left(C I^{\Theta}\right)$

## Answer: A

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13. Which of the following is the strongest acid?
A. $\mathrm{CIO}_{3}(\mathrm{OH})$
B. $\mathrm{CIO}_{2}(\mathrm{OH})$
C. $\mathrm{SO}(\mathrm{OH})_{2}$
D. $\mathrm{SO}_{2}(\mathrm{OH})_{2}$

## Answer: A

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14. Amongst the following hydroxides, the one which has the lowest value of $K_{s p}$ is:
A. $\mathrm{Mg}(\mathrm{OH})_{2}$
B. $\mathrm{Ca}(\mathrm{OH})_{2}$
C. $\mathrm{Ba}(\mathrm{OH})_{2}$
D. $\mathrm{Be}(\mathrm{OH})_{2}$
15. Which of the following solutions will have pH close to 1.0 ?
A. 100 mL of $(\mathrm{M} / 10) \mathrm{HCI}+100 \mathrm{~mL}$ of $(\mathrm{M} / 10) \mathrm{NaOH}$
B. 55 mL of $(\mathrm{M} / 10) \mathrm{HCI}+45 \mathrm{~mL}$ of $(\mathrm{M} / 10) \mathrm{NaOH}$
C. 75 mL of $(\mathrm{M} / 10) \mathrm{HCI}+90 \mathrm{~mL}$ of $(\mathrm{M} / 10) \mathrm{NaOH}$
D. 75 mL of $(\mathrm{M} / 5) \mathrm{HCI}+25 \mathrm{mLof}(\mathrm{M} / / 5) \mathrm{NaOH}{ }^{`}$

## Answer: D

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16. The pH of 0.1 M solution of the following salts increases in the order.
A. $\mathrm{NaCI}<\mathrm{NH}_{4} \mathrm{CI}<\mathrm{NaCN}<\mathrm{HCI}$
B. $\mathrm{HCI}<\mathrm{NH}_{4} \mathrm{CI}<\mathrm{NaCI}<\mathrm{NaCN}$
C. $\mathrm{NaCN}<\mathrm{NH}_{4} \mathrm{CI}<\mathrm{NaCI}<\mathrm{HCI}$
D. $\mathrm{HCI}<\mathrm{NaCI}<\mathrm{NaCN}<\mathrm{NH}_{4} \mathrm{CI}$

## Answer: B

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17. For a sparingly soluble salt $A_{p} B_{q}$, the relationship of its solubility product $\left(K_{s}\right)$ with its solubility $(\mathrm{S})$ is
A. $L_{S}=S_{p+q} p_{p} q_{q}$
B. $L_{S}=S_{p+q^{2}} p_{q} q_{p}$
C. $L_{S}=S_{p q} p_{q} q_{p}$
D. $L_{S}=S_{p q}(p q)^{(p+q)}$

## Answer: A

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18. Which of the following acids has the smallest dissociation constant?
A. $\mathrm{CH}_{3} \mathrm{CHFCOOH}$
B. $\mathrm{FCH}_{2} \mathrm{CH}_{2} \mathrm{COOH}$
C. $\mathrm{B}_{1} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{COOH}$
D. $\mathrm{CH}_{3} \mathrm{CHBrCOOH}$

## Answer: C

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19. A solution containing $\mathrm{Mn}^{2+}, \mathrm{Fe}^{2+}, \mathrm{Zn}^{2+}$ and $\mathrm{Hg}^{2+}$ with a molar concentration of $10^{-3} \mathrm{M}$ each is treated with $10^{-16} \mathrm{M}$ sulphide ion solution. Which ion will precipitate first if $K_{s p}$ of $\mathrm{MnS}, \mathrm{FeS}, \mathrm{ZnS}$ and HgS are $10^{-15}, 1-^{-23}, 10^{-20}$ and $10^{-54}$ respectively ?
A. FeS
B. $M g S$
C. HgS
D. ZnS

## Answer: C

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20. A weak acid $\mathrm{HX}\left(K_{a}=10^{-5}\right)$ on reaction with NaOH gives NaX . For 0.1 M aqueous solution of NaX , the \% hydrolysis is
A. $0.01 \%$
B. 0.001 \%
C. 0.1 \%
D. 0.5 \%

## Answer: A

21. 0.1 mol of $\mathrm{RNH}_{2}\left(K_{b}=5 \times 10^{-4}\right)$ is mixed with 0.08 mol of $\mathrm{HC1}$ and diluted to $1 L$. Calculate the $\left[H^{\oplus}\right]$ in the solution.
A. $8 \times 10^{-2} M$
B. $8 \times 10^{-11} M$
C. $1.6 \times 10^{-11} M$
D. $8 \times 10^{-5} \mathrm{M}$

## Answer: B

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22. 

$$
\mathrm{Ag}^{+}+\mathrm{NH}_{3} \Leftrightarrow\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)\right]^{+} K_{1}=3.5 \times 10^{-3}
$$

$\left.\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)\right]^{+}+\mathrm{NH}_{3} \Leftrightarrow\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{+} \mathrm{K}_{2}=1.8 \times 10^{-3}:\right\}$
then, the overall formation constant of $\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{+}$is :
A. $6.08 \times 10^{-6}$
B. $6.08 \times 10^{6}$
C. $6.08 \times 10^{-9}$
D. None of these

## Answer: A

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23. The solubility product constant $\left(K_{s p}\right)$ of salts of types $M X, M X_{2}$, and $M_{3} X$ at temperature $T$ are $4.0 \times 10^{-8}, 3.2 \times 10^{-14}$, and $2.7 \times 10^{-15}$, respectively. The solubilities of the salts at temperature $T$ are in the order
A. $M X>M X_{2}>M_{3} X$
B. $M_{3} X>M X_{2}>M X$
C. $M X_{2}>M_{3} X>M X$
D. $M X>M_{3} X>M X_{2}$

## Answer: D

24. 2.5mLof $2 / 5 M$ weak mono-acidic base $\left(K_{b}=1 \times 10^{-12}\right.$ at $\left.25^{\circ} \mathrm{C}\right)$ is titrated with $2 / 15 M H C I$ in water at $25^{\circ} \mathrm{C}$. Find the concentration of $H^{\oplus}$ ions at equivalence point. $\left(K_{w}=1 \times 10^{-14} a t 25^{\circ} \mathrm{C}\right)$
a. $3.7 \times 10^{-13} \mathrm{M}$ b. $3.2 \times 10^{-7} \mathrm{M}$
c. $3.2 \times 10^{-2} M$ d. $2.7 \times 10^{-2} M$
A. $3.7 \times 10^{-13} M$
B. $3.2 \times 10^{-7} \mathrm{M}$
C. $3.2 \times 10^{-2} \mathrm{M}$
D. $2.7 \times 10^{-2} \mathrm{M}$

## Answer: D

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25. In an acidified aqueous solution of $\mathrm{Mn}^{2+}, \mathrm{Ni}^{2+}, \mathrm{Cu}^{2+}$ and $\mathrm{hg}^{2+}$ ions, $\mathrm{H}_{2} \mathrm{~S}$ gas was passed. Precipitates are
A. CuS and HgS
B. MnS and CuS
C. MnS and NiS
D. NiS and HgS

## Answer: A

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## Archives Integer

1. The dissociation constant of a substitued benzoic acid at $25^{\circ} \mathrm{C}$ is
$1.0 \times 10^{-4}$. Find the $p H$ of a 0.01 M solution of its sodium salt.

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2. Amongst the following, the total number of compounds whose aqueous solution turns red litmus paper blue is:

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3. Find the total number of diprotic acids among the following:
$\mathrm{H}_{3} \mathrm{PO}_{4}, \mathrm{H}_{2} \mathrm{SO}_{4}, \mathrm{H}_{3} \mathrm{PO}_{3}$

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4. In 1 L saturated solution of $\mathrm{AgCl}\left[K_{S P}(\mathrm{AgCl})=1.6 \times 10^{-10}\right], 0.1$ mole of $\mathrm{CuCl}\left[K_{S P}(\mathrm{CuCl})=1.0 \times 10^{-6}\right]$ is added. The resulrant concentration of $\mathrm{Ag}^{+}$in the solution is $1.6 \times 10^{-x}$. The value of x is:

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Archives Fill In The Blanks

1. The conjugate base of $\mathrm{HSO}_{4}^{-}$in aqueous solution is $\qquad$
2. An element which can exist as a positive ion in acidic solution and also as a negative ion in basic solution is said to be.....

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3. Silver chloride is sparingly soluble in water because its lattice energy is greater than $\qquad$

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4. $\left(\mathrm{CH}_{3}(\mathrm{OH})_{2}^{+}\right)$is...acidic than $\left(\mathrm{CH}_{3} \mathrm{NH}_{3}^{+}\right)$.

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5. In the reaction $I^{-}+I_{2} \rightarrow I_{3}^{-}$, the Lewis acid is

## Archives True/False

1. Aluminium chloride $\left(\mathrm{AlCl}_{3}\right)$ is a Lewis acid because it can donate electrons.

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2. The solubility of sodium hydroxide increases with increase of temperature.

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3. Give the decreasing order of the acidic properties of oxides.
a. ZnO , b. $\mathrm{KO}_{2}$, c
c. $\mathrm{P}_{2} \mathrm{O}_{5}$,
d. MgO

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## Archives Subjective

1. How much moles of sodium propionate should be added to $1 L$ of an aqueous solution containing 0.020 mol of propionic acid to obtain a buffer solution of $p H 4.75$ ? What will be the $p H$ if 0.010 mol of HCI is dissolved in the above buffer solution. Compare the last $p H$ value with the pHof0.010MHCI solution. Dissociation constant of propionic acid, $K_{a}$, at $25^{\circ} \mathrm{C}$ is $1.34 \times 10^{-5}$.

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2. Given reason for the statement that the pH of an aqueous solution of sodium acetate is more than 7.

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3. 20 mL of 0.2 M sodium hydroxide is added to 50 mL of 0.2 Macetic acid to give 70 mL of the solution. What is the pH of this solution. Calculate the
additional volume of 0.2 MNaOh required to make the pH of the solution
4.74. (Ionisation constant of $\mathrm{CH}_{3} \mathrm{COOh}$ is $1.8 \times 10^{-5}$ )

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4. $K_{a}$ for $H A$ is $4.9 \times 10^{-8}$. After making the necessary approximation, calculate for its decimolar solution,

## $\Theta$

a. \% dissociation b. OH concentration
c. $p H$

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5. A solution contains a mixture of $\mathrm{Ag}^{+}(0.10 \mathrm{M})$ and $\mathrm{Hg}_{2}^{2+}(0.10 \mathrm{M})$ which are to be separated by selective precipitation. Calculate the maximum concentration of iodide ion at which one of them gets precipitated almost completely. $\quad\left(K_{S P} O f A g I=8.5 \times 10^{-17} \quad\right.$ and $\quad K_{S P} \quad$ of $H g_{2} I_{2}=2.5 \times 10^{-26}$ )
6. The concentration of hydrogen ions in a $0.2 M$ solution of formic acid is $6.4 \times 10^{-3} \mathrm{molL}^{-1}$. To this solution, sodium formate is added so as to adjust the concentration of sodium formate to $1 \mathrm{molL}^{-1}$. What will be the pH of this solution? The dissociation constant of formic acid is $2.4 \times 10^{-4}$ and the degree of dissociation fo sodium formate is 0.75 .

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7. The solubility of $\mathrm{Mg}(\mathrm{OH})_{2}$ in pure water is $9.57 \times 10^{-3} \mathrm{gL}$. . Calculate its solubility I $\left(\mathrm{gL}^{-1}\right)$ in $0.02 \mathrm{M} \mathrm{Mg}\left(\mathrm{NO}_{3}\right)_{2}$ solutions.

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8. What is the pH of the solution when 0.20 mol of HCI is added to 1 L of a solution containing
a. 1 M each of acetic acid and acetate ion.
b. 0.1 Meach of aceta acid and acetate ion.

Assume the total volume is $1 L . K_{a}$ for acetic acid is $1.8 \times 10^{-5}$.

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9. How many moles of HCl will be required to preapare one litre of a buffer solution (containing NaCN and HCN ) of pH 8.5 using 0.01 gram formula mass of $\mathrm{NaCN} . K_{a}$ for $\mathrm{HCN}=4.1 \times 10^{-10}$ ?

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10. Freshly prepared aluminium and magnesium hydroxides are stirred vigorously in a buffer solution containing 0.25 M of ammonium chloride and $0.05 M$ of ammonium hydroxide. Calculate the concentration of aluminium and magesium ions in solution $\left(\mathrm{K}_{b} \mathrm{NH}_{4} \mathrm{OH}=1.8 \times 10^{-5}, K_{s p} \mathrm{Mg}(\mathrm{OH})_{2}=6 \times 10^{-10}, K_{\text {sp }} \mathrm{Al}(\mathrm{OH})_{3}=6 \times 10^{-32}\right.$
11. What is the $p H$ of $1 M$ solution of acetic acid. To what volume one litre of this solution be diluted so that pH of the resulting solution will be twice of the original value. $K_{a}=1.8 \times 10^{-5}$

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12. A 50 mL solution of weak base BOH is titrated with 0.1 NHCI solution. The pH of solution is found to be 10.04 and 9.14 after the addition of 5.0 mL and 20.0 mL of acid respectively. Find out $K_{b}$ for weak base.

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13. The $K_{S P}$ of $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ at $25^{\circ} \mathrm{C}$ is $1.29 \times 10^{-11} \mathrm{~mol}^{3} \mathrm{~L}^{-3}$. A solution of $\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ containing 0.152 mole in 500 mL water is shaken at $25^{\circ} \mathrm{C}$ with excess of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$ till the equilbirium is reached.
$\mathrm{Ag}_{2} \mathrm{CO}_{3}+\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4} \Leftrightarrow \mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}+\mathrm{K}_{2} \mathrm{CO}_{3}$
Ar equilibrium the solution contains 0.0358 mole of $\mathrm{K}_{2} \mathrm{CO}_{3}$. Assuming
degree of dissociation of $\mathrm{K}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ and $\mathrm{K}_{2} \mathrm{CO}_{3}$ to be same, calculate $K_{S P}$ of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$.

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14. The solubility product $K_{\text {sp }}$ of $\mathrm{Ca}(\mathrm{OH})_{2}$ at $25^{\circ} \mathrm{C}$ is $4.42 \times 10^{-5} \mathrm{~A} 500$ mL of saturated solution of $\mathrm{Ca}(\mathrm{OH})_{2}$ is mixed with equal volume of 0.4 M NaOH . How much $\mathrm{Ca}(\mathrm{OH})_{2}$ in milligrams is precipitated?

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15. The $p h$ of blood stream is maintained by a proper balance of $\mathrm{H}_{2} \mathrm{CO}_{3}$ and $\mathrm{NaHCO}_{3}$ concentrations. What volume of $5 \mathrm{MNaHCO}_{3}$ solution, should be mixed with 10 mL sample of blood which is 2 M in $\mathrm{H}_{2} \mathrm{CO}_{3}$ in order to maintain a pH of 7.4. $\mathrm{K}_{a}$ for $\mathrm{H}_{2} \mathrm{CO}_{3}$ in blood is $7.8 \times 10^{-7}$ ?

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16. An aqueous solution of a metal bromode $\mathrm{MBr}_{2}(0.05 M)$ is saturated with $\mathrm{H}_{2} \mathrm{~S}$. What is the minimum pH at which MS will precipitate? $\mathrm{K}_{s p}$ for $M S=6.0 \times 10^{-21}$. Concentration of satured $H_{2} S=0.1 M, K_{1}=10^{-7}$ and $K_{2}=1.3 \times 10^{-13}$ for $H_{2} S$.

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17. For the reaction
$A g(C N)_{2}^{\ominus} \Leftrightarrow A g^{\oplus}+2 C N^{\ominus}$, the $K_{c}$ at $25^{\circ} \mathrm{C}$ is $4 \times 10^{-19}$ Calculate $\left[A g^{\oplus}\right]$ in solution which was originally 0.1 M in KCN and 0.03 M in $\mathrm{AgNO}_{3}$.

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18. Calculate the pH of an aqueous solution of 1.0 M ammonium formate assuming complete dissociation. $\left(p K_{a}\right.$ of formic acid $=3.8$ and $p K_{b}$ of ammonia $=4.8$ )

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19. What is the pH of a 0.50 M aqueous NaCN solution ? $\left(p K_{b} \circ f C N^{-}=4.70\right)$

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$\oplus$
20. The ionization constant of $\mathrm{NH}_{4}$ ion in water is $5.6 \times 10^{-10}$ at $25^{\circ} \mathrm{C}$. $\oplus \quad \ominus$
The rate constant the reaction of $\mathrm{NH}_{4}$ and OH ion to form $\mathrm{NH}_{3}$ and $\mathrm{H}_{2} \mathrm{O}$ at $25^{\circ} \mathrm{C}$ is $3.4 \times 10^{10} \mathrm{Lmol}^{-1} \mathrm{~S}^{-1}$. Calculate the rate constant for proton transfer from water to $\mathrm{NH}_{3}$.

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21. A sample of AgCl was treated with 5.00 mL of $1.5 \mathrm{M} \mathrm{Na} a_{2} \mathrm{CO}_{3}$ solubility to give $\mathrm{Ag}_{2} \mathrm{CO}_{3}$. The remaining solution contained $0.0026 \mathrm{gofCI}^{-}$per litre.

Calculate the solubility product of AgCl . $\left(K_{S P} f\right.$ or $\left.\mathrm{Ag}_{2} \mathrm{CO}_{3}=8.2 \times 10^{-12}\right)$

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22. An acid type indicator, H In differs in colour from its conjugate base $\left(I n^{-}\right)$. The human eye is sensitive to colour differences only when the ratio $\left[\mathrm{In}^{-}\right] /[\mathrm{HIn}]$ is greater than 10 or smaller than 0.1 . What should to observe a complete colour change ? $\left(K_{a}=1.0 \times 10^{-5}\right)$

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23. What will be the resultant pH , when 200 mL of an aqueous solution of $\operatorname{HCI}(p H=2.0)$ is mixed with 300 mL of an aqueous solution of $\mathrm{NaOH}(\mathrm{pH}=12.0) ?$

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24. Given: $\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{+} \Leftrightarrow \mathrm{Ag}^{+} 2 \mathrm{NH}_{3}, \mathrm{~K}_{\mathrm{C}}=6.2 \times 10^{-8}$ and $K_{S P}$ of $\mathrm{AgCI}=1.8 \times 10^{-10}$ at 298 K . Calculate the concentration of the complex in 1.0 M aqueous ammonia.
25. The solubility of $\mathrm{Pb}(\mathrm{OH})_{2}$ in water is $6.7 \times 10^{-6} \mathrm{M}$. Calculate the solubility of $\mathrm{Pb}(\mathrm{OH})_{2}$ in a buffer solution of $\mathrm{pH}=8$.

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26. The average concentration of $\mathrm{SO}_{2}$ in the atmosphere over a city on a cetrain day is 10 ppm , when the average temperature is 298 K . Given that the solubility of $\mathrm{SO}_{2}$ in water at 298 K is 1.3653 mol litre $^{-1}$ and the $p K_{a}$ of $\mathrm{H}_{2} \mathrm{SO}_{3}$ is 1.92 , estimate the pH of rain on that day.

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27. 500 mL of 0.2 M aqueous solution of acetic acid is mixed with 500 mL of 0.2 HCI at $25^{\circ} \mathrm{C}$.
a. Calculate the degree of dissociation of acetic acid in the resulting solution and pH of the folution.
b. If 6 g of NaOH is added to the above solution determine the final pH. $\left[\mathrm{K}_{a}\right.$ of $\mathrm{CH}_{3} \mathrm{COOH}=2 \times 10^{-5}$.

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28. 0.1 MNaOH is titrated with 0.1 MHA till the end point. $K_{a}$ of HA is $5.6 \times 10^{-6}$ and degree of hydrolysis is less compared to 1 . Calculate pH of the resulting solution at the end point ?

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