

**Daily Tutorial Sheet-2** 

JEE Main (Archive)

**16.(D)** pH = 1

$$\therefore$$
 [H<sup>+</sup>] = 10<sup>-1</sup> = 0.1 M

$$pH = 2$$

$$\therefore$$
 [H<sup>+</sup>] = 10<sup>-2</sup> = 0.01 M

For dilution of HCl,  $M_1V_1 = M_2V_2$ 

$$0.1\!\times\!1=0.01\!\times\!V_2$$

$$V_2 = 10 L$$

Volume of water to be added 10-1=9L

**17.(C)**  $HQ = H^+ + Q^-$ 

 $[H^+] = \sqrt{K_a C}$  by Ostwald's dilution law

$$[H^+] = 10^{-pH} = 10^{-3}M$$

$$C = 0.1 M$$

Thus, 
$$10^{-3} = \sqrt{K_a \times 0.1}$$

$$10^{-6} = K_a \times 0.1$$

$$K_a = 10^{-5}$$

## **Alternation Method**

$$HQ \rightleftharpoons H^+ + Q^-$$

Given, pH = 3. This suggests  $[H^+] = 10^{-3} \text{ M}$  at equilibrium =  $10^{-3} \text{ M}$ .

Hence, 
$$[HQ] = 0.1 \text{ M} - 10^{-3} \text{ M} = 0.1 \text{ M} [10^{-3} \text{ M} < << 0.1 \text{ M}]$$

 $K_a$  for the above reaction is given by:

$${\rm K_a} = \frac{{{{[{\rm{H^+}}]}\left[ {{{\rm{Q}}^ - }} \right]}}}{{{{[{\rm{HQ}}]}}}} = \frac{{{{[{\rm{10}}^{ - 3}]}\left[ {{\rm{10}}^{ - 3}} \right]}}}{{{{[{\rm{0.1}}]}}}}$$

$$K_a = 1 \times 10^{-5}$$

**18.(C)**  $H_2A \rightleftharpoons HA^- + H^+$ 

$$\therefore \qquad \quad K_1 = 1.0 \times 10^{-5} = \frac{[H^+][HA^-]}{[H_2A]} \, (given)$$

$$HA^- \rightarrow H^+ + A^{2-}$$

$$K_2 = 5.0 \times 10^{-10} = \frac{[H^+][A^{2-}]}{[HA^-]}$$
(given)

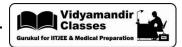
$$K = \frac{[H^+]^2[A^{2-}]}{[H_2A]} = K_1 \times K_2 = (1.0 \times 10^{-5}) \times (5 \times 10^{-10}) = 5 \times 10^{-15}$$

**19.(C)** Let molar solubility of  $Cr(OH)_3 = S \text{ mol } L^{-1}$ 

$$\operatorname{Cr}(\operatorname{OH})_3(S) \rightleftharpoons \operatorname{Cr}_s^{3+}(\operatorname{aq}) + 3\operatorname{OH}_{3s}^-(\operatorname{aq}) \lim_{x \to \infty}$$

$$\rm K_{sp} = 1.6 \times 10^{-30} = [Cr^{3+}] \ [OH^-]^3 = (S)(3S)^3 = 27 \ S^4$$

$$\therefore \qquad S^4 = \frac{1.6 \times 10^{-30}}{27} \quad \therefore \quad S = \sqrt[4]{\frac{1.6 \times 10^{-30}}{27}}$$



- **20.(C)** The pK<sub>a</sub> values of  $HCO_3^-$ ,  $H_3O^+$  and  $HSO_4^-$  are 10.25, -1.74 and 1.92 respectively and  $HSO_3$  F is super acid. (iv) > (ii) > (ii) > (i)
- **21.(A)** pH of a solution = 5

$$\therefore$$
  $[H^+] = 10^{-pH} = 10^{-5}$ 

HA (a weak acid) ionizes as

$$HA \implies H^+ + A^-$$
  
1.0 0 0  
 $(1-10^{-5}) 10^{-5} 10^{-5}$ 

$$K_{a} \frac{[H^{-}][A^{-}]}{[HA]} = \frac{10^{-5} \times 10^{-5}}{(1 - 10^{-5})} = \frac{10^{-10}}{1} [1 >>> 10^{-5}] = 1 \times 10^{-10} M$$

- **22.(A)** Only in reaction (II), H<sub>2</sub>PO<sub>4</sub><sup>-</sup>, gives H<sup>+</sup> to H<sub>2</sub>O. Thus, behaves as an acid.
- **23.(C)**  $H_2CO_3 \rightleftharpoons H^+ + HCO_3^-; K_1 = 4.2 \times 10^{-7}$

$$HCO_3^- \rightleftharpoons H^+ + CO_3^{2-}$$
;  $K_2 = 4.8 \times 10^{-11}$ 

$$K_1 >> K_2$$

$$\therefore \qquad [H^+] = [HCO_3^-]$$

$$K_2 = \frac{[H^+][CO_3^{2-}]}{[HCO_3^-]}$$

So, 
$$[CO_3^{2-}] = K_2 = 4.8 \times 10^{-11}$$

**24. (B)**  $[AgBr] = [Ag^+] = 0.05 M$ 

$$K_{sp}[AgBr] = [Ag^+][Br^-]$$

$$\Rightarrow \qquad [Br^{-}] = \frac{K_{sp}(AgBr)}{[Ag^{+}]} = \frac{5.0 \times 10^{-13}}{0.05} = 10^{-11} M \text{ [mol L}^{-1]}$$

Moles of KBr needed to precipitate AgBr = [Br $^-$ ]  $\times$  V =  $10^{-11}$  mol L $^-$ 1  $\times$ 1 L =  $10^{-11}$  mol

Therefore, amount of KBr needed to precipitate  $AgBr = 10^{-11} mol \times 120 g mol^{-1} = 1.2 \times 10^{-9} g$ 

**25. (B)**  $Mg(OH)_2 \rightleftharpoons Mg^{2+} + 2OH^{-}$ 

$$K_{sp} = [Mg^{2+}][OH^{-}]^{2}$$

$$\text{[OH^-]} = \sqrt{\frac{K_{sp}}{[Mg^{2+}]}} = \sqrt{\frac{1 \times 10^{-11}}{0.001}} = 10^{-4}$$

$$pOH = -log [OH^{-}] = -log [10^{-4}]$$

$$pOH = 4$$
 :  $pH = 14 - pOH = 14 - 4 = 10$ 

**26.(D)** 
$$pH = 1$$
  $\therefore$   $[H^+] = 10^{-1} = 0.1 M$ 

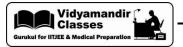
$$pH = 2$$
 :  $[H^+] = 10^{-2} = 0.01M$ 

For dilution of HCl,  $M_1V_1 = M_2V_2$ 

$$0.1 \times 1 = 0.01 \times V_2$$

$$V_2 = 10L$$

Volume of water to be added = 10 - 1 = 9L



**27. (B)** 
$$Na_2CO_3 \rightleftharpoons 2Na^+ + CO_3^{2-}$$

$$[Na_2CO_3] = [CO_3^{2-}] = 1 \times 10^{-4} M$$

$$BaCO_3 \rightleftharpoons Ba^{2+} + CO_3^{2-}$$

$$K_{sp(BaCO_3)} = [Ba^{2+}][CO_3^{2-}]$$

$$[{\rm Ba}^{2+}] = \frac{{\rm K_{sp}(BaCO_3)}}{[{\rm CO_3^{2-}}]}$$

$$[Ba^{2+}] = \frac{5.1 \times 10^{-9}}{1 \times 10^{-4}}$$

$$[Ba^{2+}] = 5.1 \times 10^{-5} M$$

Hence, at  $5.1 \times 10^{-5}$  concentration of Ba<sup>2+</sup>, a precipitate will begin to form.

**28.(C)** pH of a salt made up of weak acid and weak base is calculated by using expression

$$pH = 7 + \frac{1}{2}pK_a - \frac{1}{2}pK_b = 7 + \frac{3.2}{2} - \frac{3.4}{2} = 6.9$$

29.(D) CH<sub>3</sub>COOK is a salt of a weak acid and a strong base

. Most basic

**30.(A)** Methyl orange is used for titration of strong acid and weak base.

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