

## **Equilibrium**

**Q.No.1:** Which of the following salts is the most basic in aqueous solution?

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- A. FeCl<sub>3</sub>
- **B.** Pb(CH<sub>3</sub>COO)<sub>2</sub>
- C.  $AI(CN)_3$
- **D.** CH<sub>3</sub>COOK

**Q.No.2:** An alkali is titrated against an acid with methyl orange as indicator, which of the following is a correct combination?

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- A. Base Acid End point
  Weak Strong Yellow to pinkish red
- B. Base Acid End point
  Strong Strong Pink to colourless
- C. Base Acid End point

  Weak Strong Colourless to pink
- **D. Base Acid End point**Strong Strong Pinkish red to yellow

**Q.No.3:** An aqueous solution contains 0.10 M  $H_2S$  and 0.20 M HCl. If the equilibrium constants for the formation of  $HS^-$  from  $H_2S$  is  $1.0 \times 10^{-7}$  and that of  $S^{2-}$  from  $HS^-$  ions is  $1.2 \times 10^{-13}$  then the concentration of  $S^{2-}$  ions in aqueous solution is :

- **A.**  $6 \times 10^{-21}$
- **B.**  $5 \times 10^{-19}$
- **C.**  $5 \times 10^{-8}$
- **D.**  $3 \times 10^{-20}$

**Q.No.4:** 20 mL of 0.1 M  $H_2SO_4$  solution is added to 30 mL of 0.2 M  $NH_4OH$  solution. The pH of the resultant mixture is: [p $k_b$  of  $NH_4OH = 4.7$ ]. **JEE 2019** 

- **A.** 5.2
- **B.** 9.0
- **C.** 5.0
- **D.** 9.4

Q.No.5: Consider the following reversible chemical reactions:

$$A_2(g) + B_2(g) \xrightarrow{K_1} 2AB(g) \dots$$
 (1)

$$6AB(g) \xleftarrow{K_2} 3A_2(g) + 3B_2(g) \dots \quad (2)$$

The relation between  $K_1$  and  $K_2$  is:

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- **A.**  $K_1K_2 = \frac{1}{3}$
- **B.**  $K_2 = K_1^3$
- **C.**  $K_2 = K_1^{-3}$
- D.  $K_1K_2=3$

**Q.No.6:** The values of  $K_p/K_c$  for the following reactions at 300 K are, respectively:

(At 300 K,  $RT = 24.62 \text{ dm}^3 \text{ atm mol}^{-1}$ )

$$N_2(g) + O_2(g) = 2NO(g)$$

 $N_2O_4(g) = 2NO_2(g)$ 

$$N_2(g) + 3H_2(g) = 2NH_3(g)$$

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- **A.** 1, 24.62 dm $^3$  atm mol $^{-1}$ , 606.0 dm $^6$  atm $^2$  mol $^{-2}$
- **B.** 1, 24.62 dm<sup>3</sup> atm mol<sup>-1</sup>,  $1.65 \times 10^{-3}$  dm<sup>-6</sup> atm<sup>-2</sup> mol<sup>2</sup>
- **C.**  $1, 4.1 \times 10^{-2} \text{ dm}^{-3} \text{ atm}^{-1} \text{ mol}, 606 \text{ dm}^{6} \text{ atm}^{2} \text{ mol}^{-2}$
- **D.** 24.62 dm $^3$  atm mol $^{-1}$ , 606.0 dm $^6$  atm $^2$  mol $^{-2}$ , 1.65 × 10 $^{-3}$  dm $^{-6}$  atm $^{-2}$  mol $^2$

**Q.No.7:** 5.1 g NH<sub>4</sub>SH is introduced in 3.0 L evacuated flask at 327°C. 30% of the solid NH<sub>4</sub>SH decomposed to NH<sub>3</sub> and H<sub>2</sub>S as gases. The  $K_p$  of the reaction at 327°C is (R = 0.082 L atm mol<sup>-1</sup>  $K^{-1}$ , Molar mass of S = 32 g mol<sup>-1</sup>, molar mass of N = 14 g mol<sup>-1</sup>)

**A.** 
$$0.242 \times 10^{-4} \text{ atm}^2$$

- **B.**  $1 \times 10^{-4} \text{ atm}^2$
- **C.**  $4.9 \times 10^{-3} \text{ atm}^2$
- **D.** 0.242 atm<sup>2</sup>

**Q.No.8:** For an elementary chemical reaction,  $A_2 \overset{k_1}{\underset{k_{-1}}{\rightleftharpoons}} 2A$ , the expression for

 $\frac{d[{
m A}]}{dt}$  is:

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- **A.**  $k_1[A_2] k_{-1}[A]^2$
- **B.**  $2k_1[A_2] k_{-1}[A]^2$
- **C.**  $k_1[A_2] + k_{-1}[A]^2$
- **D.**  $2k_1[A_2] 2k_{-1}[A]^2$

**Q.No.9:** For the equilibrium,  $2H_2O = H_3O^+ + OH^-$ , the value of  $\Delta G^\circ$  at 298 K is approximately:

- **A.**  $100 \text{ kJ mol}^{-1}$
- **B.**  $-80 \text{ kJ mol}^{-1}$
- **C.** 80 kJ mol $^{-1}$
- **D.**  $-100 \text{ kJ mol}^{-1}$

Q.No.10: Consider the reaction

$$N_2(g) + 3H_2(g) = 2NH_3(g)$$

The equilibrium constant of the above reaction is  $K_{\rm p}$ . If pure ammonia is left to dissociate, the partial pressure of ammonia at equilibrium is given by (Assume that  $p_{\rm NH_3} \ll p_{\rm total}$  at equilibrium)

- **A.**  $\frac{3^{\frac{3}{2}}K_{\rm p}^{\frac{1}{2}}P^2}{16}$
- **B.**  $\frac{K_{\rm p}^{\frac{1}{2}}P^2}{16}$
- **C.**  $\frac{K_{\rm p}^{\frac{1}{2}}P^2}{4}$
- $\mathbf{D.} \ \frac{3^{\frac{3}{2}} K_{\mathrm{p}}^{\frac{1}{2}} P^2}{4}$