

Chemical and Ionic Equilibrium Formula Sheet (JEE Main & Advanced)

1. Chemical Equilibrium Basic Formulas

Equilibrium Constant

- For reaction: $aA + bB \rightleftharpoons cC + dD$

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

K_p-K_c Relationship

$$K_p = K_c(RT)^{\Delta n}$$

where $\Delta n = (c + d) - (a + b)$

Reaction Quotient (Q)

$$Q = \frac{[C]^c[D]^d}{[A]^a[B]^b} \text{ (at any instant)}$$

Free Energy Relationships

$$\Delta G = \Delta G^\circ + RT \ln Q$$

$$\Delta G^\circ = -RT \ln K_c = -2.303RT \log K_c$$

At equilibrium: $\Delta G = 0$, $Q = K_c$

Temperature Effect (van't Hoff Equation)

$$\log(K_2/K_1) = \frac{[\Delta H^\circ/2.303R][T_2 - T_1/T_1 T_2]}$$

2. Modified Equilibrium Constants

For Reversed Reaction

$$K' = 1/K$$

For Divided Equation

$$K' = \sqrt[n]{K} \text{ (when divided by } n\text{)}$$

For Multiplied Equation

$$K' = K^n \text{ (when multiplied by } n\text{)}$$

For Multi-step Reactions

$$K = K_1 \times K_2 \times K_3 \times \dots \times K_n$$

3. Ionic Equilibrium Formulas

Ionization Constants

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][BH^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 10^{-14} \text{ at } 298K$$



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pH Related

$$\text{pH} = -\log[\text{H}^+]$$

$$\text{pOH} = -\log[\text{OH}^-]$$

$$\text{pH} + \text{pOH} = 14 \text{ (at 298K)}$$

$$\text{pKa} = -\log K_a$$

$$\text{pKb} = -\log K_b$$

4. Salt Hydrolysis

Hydrolysis Constants

For weak acid + strong base salt:

$$K_h = K_w / K_a$$

$$\text{pH} = \frac{1}{2}[\text{pK}_w + \text{pK}_a + \log C]$$

For strong acid + weak base salt:

$$K_h = K_w / K_b$$

$$\text{pH} = -\frac{1}{2}[\text{pK}_w - \text{pK}_b + \log C]$$

For weak acid + weak base salt:

$$K_h = K_w / (K_a \times K_b)$$

$$\text{pH} = \frac{1}{2}[\text{pK}_a + \text{pK}_w - \text{pK}_b]$$

5. Solubility Product (K_{sp})

General Formula

$$K_{sp} = [\text{Cation}]^a[\text{Anion}]^b$$

For Different Types

Binary (1:1): $K_{sp} = s^2$

Ternary (1:2 or 2:1): $K_{sp} = 4s^3$

Quaternary (1:3): $K_{sp} = 27s^4$

Complex (3:2): $K_{sp} = 108s^5$

6. Buffer Solutions

Henderson-Hasselbalch Equations

For acidic buffer:

$$\text{pH} = \text{pK}_a + \log\left(\frac{[\text{Salt}]}{[\text{Acid}]}\right)$$

For basic buffer:

$$\text{pOH} = \text{pK}_b + \log\left(\frac{[\text{Base}]}{[\text{Salt}]}\right)$$

Buffer Capacity (β)

$$\beta = \frac{\Delta n}{\Delta \text{pH}}$$

7. Degree of Dissociation (α)

For Weak Electrolyte

$$\alpha = \sqrt{K_a/C} \text{ [for dilute solutions]}$$

$$\alpha = [\text{ionized}]/[\text{initial}]$$

Ostwald's Dilution Law

$$K_a = C\alpha^2/(1-\alpha)$$

For very weak electrolytes ($\alpha \ll 1$):

$$K_a = C\alpha^2$$

8. Common Ion Effect

$$[H^+] = K_a \times [HA]/[A^-]$$

$$[OH^-] = K_b \times [B]/[BH^+]$$



9. Temperature Dependence

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$d(\ln K)/dT = \Delta H^\circ/RT^2$$

Important Notes:

1. All concentrations must be in mol/L for K_c
2. All pressures must be in atm for K_p
3. Temperature must be in Kelvin
4. For dilute solutions, molarity \approx molality
5. Activity coefficients are assumed to be 1 for dilute solutions