Chemistry 223

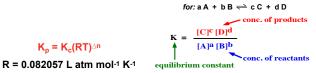


Key Equations for CH 223

This may not include all the important equations from CH 223, but most of them are included here, separated by chapter.

Good luck!

Important Equations, Constants, and Handouts from Chapter 13:



Under Any Reaction Conditions

Product concentrations Reaction quotient = $Q = \frac{[C]^c[D]^d}{[C]^d}$ [A]a[B]b

- Manipulating Equilibrium Constant Expressions
- Types of Equilibrium Constants

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

Le Chatelier's Principle MAR

Important Equations, Constants, and Handouts from Chapter 14 Part I:

 $pH_{(strong\ acid)} = - log\ C_a$ pH_(strong base) = 14 + log C_b $pH_{(weak\ acid)} = -log\ [K_a * C_a]^{1/2}$ $pH_{\text{(weak base)}} = 14 + log [K_b * C_b]^{1/2}$ $14 = pH + pOH = pK_a + pK_b$ $K_w = 1.00 * 10^{-14} = [H_3O^+][OH^-] = K_a^*K_b (25 °C)$ HOAc + H₂O ⇌ H₃O+ + OAc- $K_a = \frac{[H_3O^+][OAc^-]}{[H_3O^+][OAc^-]}$ [HOAc] $B(aq)\,+\,H_2O(\ell)\, \, {\color{red}\Longleftrightarrow}\,\, BH^+(aq)\,+\,OH^-(aq)$ $K_b = \frac{[BH^+][OH^-]}{}$

Acid-Base Theory: Brønsted theory, conjugate acid and base, strong and weak acids and bases, know the 8 strong acids and bases!, autoionization, Lewis theory, electron pair acceptor and donator, salt acidity/basicity, formation constants (K_f)

- · Manipulating Equilibrium Constant Expressions
- · Types of Equilibrium Constants
- Table of K_B and K_b values in Problem Set #2

Important Equations, Constants, and **Handouts from Chapter 14 Part II:**

Titrations (SA+SB, SB+SA, WA+SB, WB+SA) and Buffers

chapter

$$\begin{split} \mathbf{pH} &= \mathbf{pK_a} + \mathbf{log} \frac{[\mathbf{Conj.base}]}{[\mathbf{Acid}]} \\ \mathbf{pH} &= \mathbf{pK_a} + \mathbf{log} \frac{[\mathbf{mol_{Conj.base}} - \mathbf{mol_{strong.acid}})}{[\mathbf{mol_{weak.acid}} + \mathbf{mol_{strong.acid}})} \\ \mathbf{pH} &= \mathbf{pK_a} + \mathbf{log} \frac{[\mathbf{mol_{Conj.base}} + \mathbf{mol_{strong.base}})}{[\mathbf{mol_{weak.acid}} - \mathbf{mol_{strong.base}})} \end{split}$$

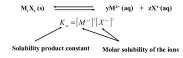
- Manipulating Equilibrium Constant Expressions
- Types of Equilibrium Constants
- Table of K_a and K_b values in Problem Set #2
- · Titration Guide

MAR · Buffers and Henderson-Hasselbalch Guide

Pre-Equivalence Region: $pH = -\log\left(\frac{n_{xx} - n_{xb}}{V_{xx} + V_{xb}}\right)$ re Region: $pH = -m_b \left\{ V_{xx} + V_{xx} \right\}$ pH = 7 $ce Region: <math>pH = 14 + log \left(\frac{n_{th} - n_{xx}}{V_{sh} + V_{ss}} \right)$ Initial Region: $pH = 14 + log \left(\frac{n_{cb} - n_{ca}}{V_{ss} + V_{sb}} \right)$ Pre-Equivalence Region: $\begin{aligned} & pH = 17 + \log \left[\frac{1}{V_{xx}} + V_{xy} \right] \\ & Equivalence: & pH = 7 \end{aligned}$ Post-Equivalence Region: $pH = -\log \left(\frac{n_{xx} - n_{xy}}{V_{xx}} + V_{xy} \right)$ $\label{eq:Weak Actid + Strong Base:} \begin{aligned} & \underline{\mathbf{Weak Actid + Strong Base:}} \\ & \underline{\mathbf{pH}} = -\log \sqrt{K_s} * \frac{\mathbf{n}_{n_s}}{V_{v_s}} \\ & \underline{\mathbf{pH}} = \underline{\mathbf{pK}}_s + \log \left(\frac{\mathbf{n}_{n_s}}{\mathbf{n}_{n_s} - \mathbf{n}_{n_s}} \right) \end{aligned}$ PFE-equivalence Region: $\begin{aligned} & \left\langle \mathbf{n}_{n_{n}} - \mathbf{n}_{n_{n}} \right\rangle \\ & Half-Equivalence Region: \\ & Equivalence: \end{aligned} \qquad \begin{aligned} & \mathbf{pH} = \mathbf{pK}_{\mathbf{x}} \\ & \mathbf{pH} - 14 + \log \sqrt{\frac{\mathbf{K}_{n}}{\mathbf{K}_{n}}} * \frac{\mathbf{n}_{n_{n}}}{\left\langle \mathbf{V}_{n_{n}} + \mathbf{V}_{n_{n}} \right\rangle} \end{aligned}$ $Poss-Equivalence \ Region: \qquad pH = 14 + log \left(\frac{n_{sb} - n_{ex}}{V_{sb} + V_{se}} \right)$ $$\label{eq:WeakBase+Strong-Acids} \begin{split} & \underline{WeakBase+Strong-Acids}: \\ & \text{Initial Region:} & pH = 14 + log \sqrt{K_s * \frac{n_{-n_s}}{V_{s,s}}} \\ & Pre\text{-}Equivalence Region:} & pH = \underline{pK_8} + log \left(\frac{n_{-n_s}-n_s}{n_{n_s}}\right) \end{split}$$ $\begin{array}{ll} \textit{Half-Equivalence Region:} & \textbf{pH} = \textbf{pK}_{a} \\ \textit{Equivalence:} & \textbf{pH} = -\log \sqrt{\frac{K_{w_{a}}}{K_{b}}} \cdot \frac{\textbf{n}_{+b}}{\left(V_{w_{0}} + V_{w_{0}}\right)} \\ \end{array}$ wivalence Region: $pH = -log \left(\frac{n_{ss} - n_{sh}}{V + V} \right)$

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Important Equations, Constants, and Handouts from Chapter 15:



· know how to predict solubility using CH 221 solubility guide

$$\begin{split} & \text{Ag*(aq)} \ + \ 2 \ \text{CN*}^{\text{1}}(\text{aq}) \ \Longleftrightarrow \ \text{Ag(CN)}_2^{\text{-1}}(\text{aq}) \\ & K_{\mathrm{f}} = \frac{\left[Ag(CN)_2^{-1}\right]}{\left[Ag^+\right]\left[CN^{-1}\right]^2} = 5.6 * 10^{18} \end{split}$$

Solubility: Common ion effect, separating salts by differences in

Handouts:

- · Types of Equilibrium Constants
- · Solubility Guide

Important Equations, Constants, and Handouts from Chapter 16:

 $\Delta H_{sys} \circ = \Sigma \Delta H \circ (products) - \Sigma \Delta H \circ (reactants)$ ΔS_{sys} ° = ΣS ° (products) - ΣS ° (reactants) $\Delta G_{sys}^{\circ} = \Sigma \Delta G^{\circ}$ (products) - $\Sigma \Delta G^{\circ}$ (reactants) $\Delta G^{\circ} = \Delta H^{\circ} - T \Delta S^{\circ}$

 $\Delta G_{rxn}^{\circ} = -RT \ln K$ R = 8.3145 J mol-1 K-1 $\Delta G_{rxn} = \Delta G_{rxn} + RT \ln Q$

Handouts:

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- · Types of Equilibrium Constants
- · Thermodynamic Values (Problem Set #5)

- · know the three laws of thermodynamics!
- know the difference between enthalpy and entropy and how they relate to Gibbs free energy
- know how to calculate enthalpy (CH 221) entropy and Gibbs energy (this chapter)
- know how the sign of ∆G relates to spontaneity (and also ∆Suniverse)

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Important Equations, Constants, and Handouts from Chapter 17:

Redox Reactions: oxidation, reduction, LEO, GER, oxidizing agent, reducing agent, anode, cathode, galvanic/voltaic cells, electrolysis (electrolytic cells), shorthand notation for galvanic cells, SHE electrode

- know how to balance redox reactions in acid or base conditions
- be able to calculate E^o and E for cells

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$$\Delta G^{\circ} = - n F E^{\circ}$$
 $Amps = \frac{coulombs}{seconds}$
 $E = E^{\circ} - (RT/nF) ln Q$
 $R = 8.3145 J mol^{-1} K^{-1}$
 $F = 9.6485 \times 10^{4} C/mol e^{-1}$

$$E^{\circ} = \frac{RT}{nF} \, \ln \, K$$

Handouts:

 Thermodynamic Values and Electrochemical Cell Values (Problem Set #5)