OTHS Pre-AP Chemistry Equations and Constants

The following symbols have the definitions specified unless otherwise noted.

g = gram(s) nm = nanometer(s) atm = atmosphere(s) L, mL = liter(s), milliliter(s)	J, kJ = joule(s), kilojoule(s) mol = mole(s) mm Hg = millimeters of mercury Pa, kPa = pascal(s), kilopascal(s)
ATOMS, ENERGY, & ELECTRONS	
$E_{photon} = h\nu$ $c = \lambda\nu$ $E_{photon} = \frac{hc}{\lambda}$	$E = energy$ $\nu = frequency$ $\lambda = wavelength$ Planck's constant, h = 6.626 × 10 ⁻³⁴ J · s Speed of light, c = 2.998 × 10 ⁸ m · s ⁻¹ Avogadro's number = 6.022 × 10 ²³ mol ⁻¹
ACIDS and BASES $K_{a} = \frac{[H^{+}][A^{-}]}{[HA]}$ $K_{b} = \frac{[OH^{-}][HB^{+}]}{[B]}$ $K_{w} = [H^{+}][OH^{-}] = 1.0 \times 10^{-14} @ 25^{\circ}C$ $K_{w} = K_{a} \times K_{b}$ $pH = -\log[H^{+}], pOH = -\log[OH^{-}]$ $14 = pH + pOH$	Equilibrium Constants K _a (weak acid) K _b (weak base) K _W (water)
GASES, LIQUIDS, AND SOLUTIONS	
$PV = nRT$ $P_{total} = P_1 + P_2 + P_3 \dots$ $\frac{P_1V_1}{T_1n_1} = \frac{P_2V_2}{T_2n_2}$ $K = \ ^\circ C + 273.15$ $n = \frac{m}{molar \ mass}$ $Kinetic \ Energy = \frac{1}{2}mv^2$ $Molarity, \qquad M = \frac{moles \ solute}{liters \ solution}$ $Dilution \ formula, \qquad M_1V_1 = M_2V_2$ $D = \frac{m}{V}$	$P = \text{pressure}$ $V = \text{volume}$ $T = \text{temperature}$ $n = \text{number of moles of gas}$ $R, \text{ Ideal Gas Constant} = 8.314 L \cdot kPa \cdot mol^{-1}K^{-1}$ $= 0.08206 L \cdot atm \cdot mol^{-1}K^{-1}$ $= 62.36 L \cdot torr \cdot mol^{-1}K^{-1}$ $= 8.314 J \cdot mol^{-1}K^{-1}$ $K = \text{Kelvin temperature}$ $1atm = 760 mm Hg = 760 torr = 101.3kPa$ $STP = 273.15 K and 1.000 atm$ $Volume of an Ideal Gas @ STP = 22.4 L \cdot mol^{-1}$ $v = \text{velocity}$ $D = \text{density}, m = \text{mass}$ $1cm^3 = 1mL$

THERMOCHEMISTRY

 $q = mc\Delta T$

$$\Delta H^{\circ} = \sum \Delta H_{f}^{\circ}(products) - \sum \Delta H_{f}^{\circ}(reactants)$$

q = heat m = mass c = specific heat capacity T = temperature

 $H^{\circ} = standard enthalpy$ $\Delta H^{\circ}_{f} = standard enthalpy of formation$

OTHER FORMULAE

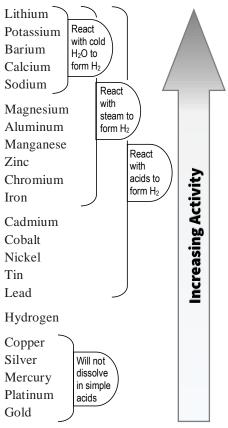
 $Percent \ error \ = \ \left| \frac{accepted \ value \ - \ experimental \ value}{accepted \ value} \right| \times 100$

Percent Yield = $\left(\frac{actual yield}{theoretical yield}\right) \times 100$

SIGNIFICANT FIGURE RULES

- 1. Non-zero digits and zeros between non-zero digits are always significant.
- 2. Leading zeros are not significant.
- 3. Trailing zeros are significant if a decimal point is shown.
- 4. In scientific notation, all digits of the coefficient are significant.
- 5. In a logarithm, there are as many digits after the decimal point as there are significant figures in the original value.

ACTIVITY SERIES FOR METALS



SOLUBILITY RULES FOR SALTS

These rules are written by priority, top to bottom. Rules higher in the list override rules lower in the list. Each statement implies that the listed ion behaves as the title states.

*Examining several sources will yield several slightly different sets of solubility rules. For our work and assessments, these are the only rules to consider.

Always Soluble

- 1. alkali metal cations (Li⁺, Na⁺, K⁺, Rb⁺, Cs⁺) and NH₄⁺
- 2. NO_3^- , $C_2H_3O_2^-$, NO_2^- , CIO_3^- , CIO_4^- , CN^- , HCO_3^-

Generally Soluble

- 3. Cl⁻, Br⁻, I⁻ except when bound to Hg_2^{2+} , Ag^+ , Pb^{2+}
- 4. SO_4^{2-} except Ca^{2+} , Sr^{2+} , Ba^{2+} , Hg_2^{2+} , Ag^+ , Pb^{2+}

Generally Insoluble

- 5. F^- , CO_3^{2-} , PO_4^{3-} , S^{2-} , SO_3^{2-} , $C_2O_4^{2-}$, CrO_4^{2-} , $Cr_2O_7^{2-}$ except Rule 1.
- 6. O²⁻, OH⁻ except Rule 1. Oxides and hydroxides of Ca²⁺, Sr²⁺, Ba²⁺ are slightly soluble (this is still considered insoluble!)