OTHS Pre-AP Chemistry Equations and Constants

The following symbols have the definitions specified unless otherwise noted.

g nm atm L, mL	<pre>= gram(s) = nanometer(s) = atmosphere(s) = liter(s), milliliter(s)</pre>	mol = mol mm Hg = mill	e(s), kilojoule(s) e(s) imeters of mercury cal(s), kilopascal(s)
ATOMS, ENER $E_{photon} = hv$ $c = \lambda v$ $E_{photon} = \frac{hc}{\lambda}$	RGY, & ELECTRONS	E = energy v = frequency $\lambda = wavelength$ Planck's constant, h Speed of light, c Avogadro's number	= $6.626 \times 10^{-34} J \cdot s$ = $2.998 \times 10^8 m \cdot s^{-1}$ = $6.022 \times 10^{23} mol^{-1}$
ACIDS and BA	SES		

$K_a = \frac{[H^+][A^-]}{[HA]}$	<u>Equilibrium Constants</u> K _a (weak acid)
$[OH^{-}][HB^{+}]$	<i>Kb</i> (weak base)
$K_b = \frac{[OH^-][HB^+]}{[B]}$	K_W (water)
$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} @ 25^{\circ}C$	
$K_w = K_a \times K_b$	
$pH = -\log[H^+], \qquad pOH = -\log[OH^-]$	
$[OH^{-}] = 10^{-pOH}, \qquad [H^{+}] = 10^{-pH}$	
14 = pH + pOH	
	1

GASES, LIQUIDS, AND SOLUTIONS

PV = nRTP = pressureV = volume $P_{total} = P_1 + P_2 + P_3 \dots$ T = temperaturen = number of moles of gas $\frac{P_1 V_1}{T_1 n_1} = \frac{P_2 V_2}{T_2 n_2}$ $= 8.314 L \cdot kPa \cdot mol^{-1}K^{-1}$ *R*, Ideal Gas Constant $= 0.08206 L \cdot atm \cdot mol^{-1}K^{-1}$ $= 62.36 L \cdot torr \cdot mol^{-1}K^{-1}$ $= 8.314 J \cdot mol^{-1}K^{-1}$ K = °C + 273.15*K* = Kelvin temperature т °C = Celsius temperature $n = \frac{1}{molar mass}$ 1atm = 760.0 mm Hg = 760.0 torr = 101.3 kPa*Kinetic Energy* $= \frac{1}{2}mv^2$ STP = 273.15 K and 1.000 atmVolume of an Ideal Gas @ STP = $22.40 L \cdot mol^{-1}$ $M = \frac{moles \ solute}{liters \ solution}$ Molarity, v = velocity Dilution formula, $M_1V_1 = M_2V_2$ D =density, m =mass $D = \frac{m}{V}$ $1cm^{3} = 1mL$

THERMOCHEMISTRY

 $q = mC\Delta T$ $\Delta H^{\circ} = \sum \Delta H_{f}^{\circ}(products) - \sum \Delta H_{f}^{\circ}(reactants)$ $C \text{ of } H_{2}O(l) = 4.184 \frac{J}{g^{\circ}C}$ $C \text{ of } H_{2}O(s) = 2.03 \frac{J}{g^{\circ}C}$ $C \text{ of } H_{2}O(g) = 2.06 \frac{J}{g^{\circ}C}$ $M^{\circ} = standard enthalpy$ $\Delta H_{f}^{\circ} = standard enthalpy \text{ of formation}$

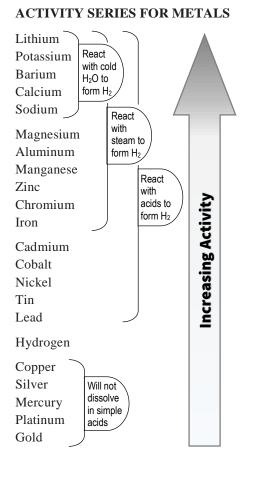
OTHER FORMULAE

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

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.



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 NH_{4^+}
- 2. NO_3^- , $C_2H_3O_2^-$, NO_2^- , ClO_3^- , ClO_4^-

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⁻, CO₃²⁻, PO₄³⁻, S²⁻, SO₃²⁻, C₂O₄²⁻, CrO₄²⁻, Cr₂O₇²⁻, CN⁻ except Rule 1.
- 6. O^{2–}, OH[–] except Rule 1. Oxides and hydroxides of Ca²⁺, Sr²⁺, Ba²⁺ are slightly soluble (this is still considered insoluble!)