General Chemistry cheat sheet DAT General Chemistry Equation Sheet

Chapter 0: G	ieneral and Lab Co	oncepts Review
Dilutions	$M_1V_1 = M_2V_2$ or $C_1V_1 = C_2V_2$	M or C = concentration V = volume
Percent Error	$\frac{(A-T)}{T} \times 100$	T = theoretical A = actual
Absorbance (Spectro- photometer)	Abs = eci	<pre>e = molar extinction coefficient (molar absorptivity) c = sample's concentration l = path length</pre>

Energy of a photon	$E_{photon} = hf = \frac{hc}{\lambda}$	h = Flanck's constant (6.63 × 10° "f' - s) f' = photon's frequency c = speed of light (8.0 × 10° "f _s) \(\lambda\) = photon's worselength
Absorption/ Emission Line Spectra	$\Delta E = E_{photom}$	-
Kinetic Energy of an electron (Photoelectric Effect)	$KE_{g-} = E_{photon} - \phi$	 φ = work function (minimum energy needed to touize electron)

Molarity	$M = \frac{moles_{solute}}{L_{solution}}$	
Molality	$m = \frac{moles_{solute}}{k q_{outpeak}}$	
Henry's Law	$P_A = k_{cc}[A]$	P _A = partial pressure of gas A k _a = Heary's Low constant (varies per problem) [All = cone, of gas A
Freezing Point Depression	$\Delta T_F = -iK_F m$	i = van't Moff factor K _p = F.P. depression constant m = molality
Boiling Point Elevation	$\Delta T_B = iK_B m$	i = van't Moff factor K _a = B.P. depression constant m = modelity
Vapor Pressure Depression (Raoult's Law)	$P_{soin} = \chi_{sois} P_{sois}^{ 0}$	P _{sobs} = VP of solution X _{sobs} = mol fract of solvent P _{sobs} = VP of pure solvent
Osmotic Pressure (rr)	$\pi = \iota MRT$	M = readerity of solute i = read t Hoff factor B = 0.002 t - other T = terms, for Kelvin

Chapter 5		
Pressure	$P = \frac{F}{A}$	F = force
	7 7	A = area
Awerage Kir Energy	netic $KE_{exp} = 3/2$	$RT = 8.314 \frac{I}{mol \cdot K}$
Root-Mean Square Spe (v)	THE PARTY	$R = 8.314 \frac{I}{\text{mol · K}}$ $M_m = \text{molar mass}$
ldeal Gas La	PV = nRT	$n = \theta$ of moles $R = 0.0021 \frac{L \cdot atm}{mol \cdot K}$
Boyle's Law	$V \propto \frac{1}{P}$	mor · K
Charles' Lav	w VecT	
Awaga deo's		
Combined C	$\frac{P_1V_1}{n_1T_1} = \frac{P_2V_2}{n_2T_2}$	
Standard Te	mo. Pel atm	*1 mol of gas - 22.41
& Pressure		at STP
Standard	All agraeous sp	ecies (it 1M
Conditions	All gaseous spe T=298 K	
Density	$\frac{P(MM)}{RT} = \frac{m}{v}$	MM = molor mass $R = 0.0021 \frac{L \cdot atm}{mol \cdot K}$ m = mass v = volume
Dalton's La Partial Pres	17. TO CO. C. C. SHOW HELD 1. 1. 10. 10. 10.	$P_B + \cdots$
Dalton's La Partial Pres	THE RESERVE THE PERSON NAMED IN COLUMN TWO IS NOT THE PERSON NAMED IN COLUMN TWO IS NAMED I	χ _s = mol fraction of gas A
Graham's L Effusion	arw of $\frac{r_1}{r_2} = \sqrt{\frac{M_{m2}}{M_{m1}}}$	r = rate of effusio M = molar mass
Real Gas Equation	aw of $\frac{r_1}{r_2} = \sqrt{\frac{M_{m2}}{M_{m3}}}$ $(P + \frac{\alpha n^2}{V^2})(V - nb) = nRT$	a & h = constants specific to each gas an' We corrects for IMFs -ab corrects for volume
Chapter 8	3: Chemical Kinetic	5
General Rate Law	$A+B \rightarrow C+D$ rate = $k[A]^m[B]^n$	k = rate constant m & n = determined experimentally
Rate Constant Units	0 order: $k = M^1 \cdot x^{-1}$ 1^{rl} order: $k = s^{-1}$ 2^{rd} order: $k = M^{-1} \cdot 3^{rd}$ order: $k = M^{-2} \cdot 3^{rd}$	k = rate constant M = molarity s= seconds

 $k = Ae^{-E_a/a\tau}$

Archenius.

Equation

k = rate constant A = unique to each run

 $E_a = act.energy$ $R = 0.314 \frac{f}{mol \cdot K}$ T = temp.in Kelvin

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G Orfield

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