

Exam 1		Exam 1 (cont)		Exam 1 (cont)		Exam 1 (cont)		
Kelvin to Celsius	$K=C + 273$	electrostatic energy	$E_{el} = Q_1Q_2/d$	Q1 and Q2: product of charges; d: distance between charges	energy (hv) of a photon used to eject electrons from a metal surface via the photoelectric effect is equal to the sum of kinetic energy of the ejected electron (Ek) and the work function (W)	hv Ek = = Ek hv + - W W	difference in energy between two quantum states	$E = hv = -2.18 \times 10^{-18} J (1/n(f)^2 - 1/n(i)^2)$
Fahrenheit to Celsius	$F=9F/5C (C) + 32F$	joule	$1 J = 1 kg \times m^2/s^2$	$1 J = 1 N \times m$	wavelength of emitted/absorbed light when an electron transitions from one quantum state to another	$1/\text{wavelength} = 1.097 \times 10^7 m^{-1} (1/n(f)^2 - 1/n(i)^2)$	energy of an electron with a given quantum state	$E_n = -2.18 \times 10^{-18} J (1/n^2)$
density	$d=m/V$ SI: $kg/m^3$ ; g/mL or $g/cm^3$ commonly used	speed, wavelength, and frequency	$c = (\text{wavelength}) (v)$	c: speed of light - $3.00 \times 10^8 m/s$ ; wavelength: in meters; frequency (v): in $s^{-1}$ or Hz	de broglie wavelength	wavelength = $h/mu$	wavelength of emitted/absorbed light	$1/\text{wavelength} = 2.18 \times 10^{-18} J/hc (1/n(f)^2 - 1/n(i)^2)$
moles to atoms and molecules	1 mole = $6.022 \times 10^{23}$ atoms or molecules	grams to atoms or molecules	atomic mass (g) = $6.022 \times 10^{23}$ molecules	formula mass (g) = $6.022 \times 10^{23}$ molecules	kinetic energy of a moving object	$E_k = 1/2 mu^2$	avagadro's number	$6.022 \times 10^{23}$ moles



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Exam 1 (cont)		Exam 2		Exam 2 (cont)		Exam 3	
Heisenberg uncertainty principle	$\Delta x \Delta p > h/4\pi$	effective nuclear charge ( $Z_{eff}$ ) $Z_{eff} = Z - \sigma$	$F = Q_1Q_2/d^2$	% ionic character	$= u$ (observed)/ $u$ (calculated) (100%)	dipole moment	$= \text{number of electrons in bonding MO} - \text{number of electrons in antibonding MO} / 2$
energy and wavelength	$E = hc/\lambda$	wavelength (nm)	$F = Q_1Q_2/d^2$	dipole moment	$u = Q/r$	dipole moment (in debye units (D)); Q: charge magnitude; r: distance between charges (bond length)	$\text{atom} = \text{sum of molar mass of desired product} / \text{sum of molar mass of reactants}$
charge of a single electron	$-1.6022 \times 10^{-19} \text{ C}$	Coulomb's law	ionic EN difference	charge magnitude	$Q = u/r$	dilution	$\% \text{ yield} = \text{actual yield} / \text{theoretical yield} (100\%)$
atomic mass units (amu)	$1 \text{ amu} = 1.66 \times 10^{-24} \text{ g}$	nonpolar (or purely covalent) EN difference	polar EN difference	formal charge	$= \text{valence electrons} - (\text{all nonbonding electrons} + 1/2 \text{ bonding electrons})$	molarity	$M = \text{moles solute} / \text{L solution}$
angstrom	$1 \text{ \AA} = 1 \times 10^{-10} \text{ m}$	% by mass of an element	nonpolar (< .5)	electronegativity	$EN = IE_1 + EA / 2$	$M_c \times c = M_d \times d$	$M_c \times c = M_d \times d$
mass of a single electron	$9.10 \times 10^{-28} \text{ g}$		covalent EN difference	coulomb charge	$1 \text{ C} = 6.242 \times 10^{18} \text{ electrons}$	$M_c \times c = M_d \times d$	$M_c \times c = M_d \times d$
mass of a proton	$1.67262 \times 10^{-24} \text{ g}$					$M_c \times c = M_d \times d$	$M_c \times c = M_d \times d$
charge-to-mass ratio of an electron	$1.76 \times 10^8 \text{ C/g}$					$M_c \times c = M_d \times d$	$M_c \times c = M_d \times d$



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Exam 4 (cont)			Exam 4 (cont)			Exam 4 (cont)			Exam 4 (cont)			
average kinetic energy of a group of gas molecules	$u^2 = \frac{uN^2}{N}$	$u^2$ : average speed for all the molecules in the sample; mean square speed	comparing Urms(1)/Urms(2) = number of molecules in different gas samples	Urms(1)/Urms(2) = square root of molar mass (2)/molar mass (1)	rate = 1/square root of molar mass	rate of diffusion or effusion is inversely proportional to the square root of the molar mass	pressure P = force/area	SI unit of force: Newton (1 N = 1kg x m/s <sup>2</sup> )	SI unit of pressure: Pa, 1 Pa = 1 N/m <sup>2</sup>	Pressure by Pascal	pressure of a fixed amount of gas at constant temperature is inversely proportional to the volume of the gas	
total kinetic energy of one mole of any gas	$E_k = \frac{3}{2} RT$	R: 8.314 J/K x mol	T: temperature in Kelvin	molar mass in kg/mol			pressure exerted by a column of fluid	P = hdg	P: pressure in Pa	h: height of column in meters	d: density of fluid in kg/m <sup>3</sup>	g: gravitational constant - 9.80665 m/s <sup>2</sup>
root-mean-square speed	$U_{rms} = \sqrt{\frac{3RT}{molar\ mass}}$	R: 8.314 J/K x mol										



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Exam 4 (cont)			Exam 4 (cont)			Exam 4 (cont)			Exam 4 (cont)				
charles law	$V_1/T_1 = V_2/T_2$	volume of a fixed amount of gas at constant pressure is directly proportional to the absolute temperature of the gas	avogadros law	$V_1/n_1 = V_2/n_2$	volume of a sample of gas at constant temperature and pressure is directly proportional to the number of moles in the sample	ideal gas equation	$PV = nRT$	R: 0.08206 L x atm/K x mol	T partial pressure	P total = sum of partial pressures	$X_i = n_i/n_{total}$	$X_i = P_i/P_{total}$	
						density of a gas	$d = P(\text{molar mass})/RT$	molar mass in kg/mol	R: amount of reactant consumed	$n = P \times (V/RT)$ at constant volume and temperature	n: number of moles consumed		
						molar mass of a gas	$dRT/P$	R: 0.08206 L x atm/K x mol	molar mass: in kg/mol	pressure	P total = P O <sub>2</sub> + P H <sub>2</sub>		
						van der waals equation	$(P + a n^2/V^2)(V - nb) = nRT$	a and b depend on the element					
			combined gas law	$P_1V_1/n_1T_1 = P_2V_2/n_2T_2$	$P_1V_1/T_1 = P_2V_2/T_2$	compressibility factor	$Z = PV/RT$						



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