

Chemistry 20-A
Dr. E.R. Scerri
Mid Term Exam
November, 2014

50 minutes

PROBLEM	SCORE	Max Score
1	14	14
2	6	12
3	16	16
4	9	12
total	45	54

(Last name first) Mortel Dominic

Signature D Mortel

ID # 904 227 174

Instructions: This exam has 4 questions plus a periodic table at end of exam. Verify you have the right number of pages before you begin. Different questions carry different numbers of points. Write your name on each page. Raise your hand if you don't understand a question. **SHOW YOUR WORK!** No credit will be given for an unsubstantiated or illegible answer. Write legibly, use proper units throughout and use significant figures in all answers. Good luck!

Possibly useful information:

$$12 \text{ inches} = 1 \text{ foot}, \quad 2.54 \text{ cm} = 1 \text{ inch.}$$

$$h = 6.63 \times 10^{-34} \text{ J sec} \quad N_o = 6.02 \times 10^{23} \text{ mol}^{-1} \quad c = 3.00 \times 10^8 \text{ m sec}^{-1}$$

$$m_e = 9.11 \times 10^{-31} \text{ kg} \quad 1 \text{ a.m.u.} = 1.66 \times 10^{-27} \text{ kg} \quad 1 \text{ \AA} = 10^{-10} \text{ m}$$

$$g = 9.81 \text{ m/s} \quad 1 \text{ eV} = 1.602 \times 10^{-19} \text{ J}$$

$$V_{\text{electrostatic}}(r) \propto Q_1 Q_2 / r; \quad \text{K.E.} = 1/2 m v^2 = p^2 / 2m$$

$$F = -\Delta V / \Delta r \quad \lambda v = c \quad \Delta E = E_f - E_i = \epsilon_{\text{photon}} = h\nu \quad h\nu = h\nu_o + \text{K. E. (electron)}$$

$$\lambda_{\text{mass}} = \frac{h}{mv} = \frac{h}{p} \quad \Delta p \Delta x \geq \frac{h}{4\pi}$$

$$\text{Probability density} = |\Psi_n(r)|^2$$

$$E_n = - (2.18 \times 10^{-18} \text{ J}) Z^2 / n^2$$

$$r_n = (0.529 \text{ \AA}) n^2 / Z_{\text{eff}}$$

$n - \ell - 1$ spherical (radial) nodes; ℓ angular nodes; $n - 1$ total nodes

$$\text{Bond order} = (\# \text{ bonding e}^- - \# \text{ antibonding e}^-) / 2$$

Spectrochemical series (abbreviated) $\text{Cl}^- > \text{F}^- > \text{H}_2\text{O} > \text{NH}_3 > \text{en} > \text{CN}^-$

1. Consider the following unbalanced equation



Assume that 40.0 g of $(\text{NH}_4)_3\text{PO}_4$ are reacted with 25.0 g of $\text{Pb(NO}_3)_4$.

How many grams of NH_4NO_3 will be produced assuming that the yield for the reaction is 68%? Assume the following approximate molecular masses for the four substances in the above reaction, from left to right, 149, 455, 991 and 80 amu. (10 points)

$$\frac{40.0 \text{ g (NH}_4\text{)}_3\text{PO}_4}{149 \text{ g}} \times \frac{1 \text{ mol}}{1 \text{ mol}} \times \frac{12 \text{ mol}}{4 \text{ mol}} = 8.05 \text{ mol}$$

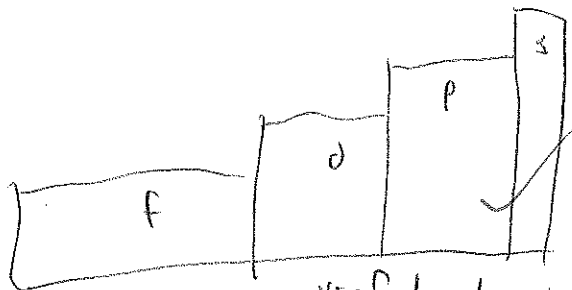
$\frac{10}{10}$

limiting reagent

$$\frac{25.0 \text{ g Pb(NO}_3\text{)}_4}{455 \text{ g}} \times \frac{1 \text{ mol}}{1 \text{ mol}} \times \frac{12 \text{ mol}}{3 \text{ mol}} = 2.20 \text{ mol NH}_4\text{NO}_3$$

$$= 17.6 \text{ g} \times 0.68 = 12.0 \text{ g NH}_4\text{NO}_3$$

1(b) Sketch a left-step periodic table and mention it's possible advantages and disadvantages (4)



This table can easily $\frac{4}{4}$ show which atoms ~~are~~ in ~~each orbital~~ have valence e^- in each orbital, however it doesn't show useful trends like orbital radius or ionization energy. Like the middle-long form can.

It also doesn't help to see the # of valence e^- if each atom

2(a). In quantum mechanics the operator for linear momentum in the x direction is given by

$$p_x = -i \hbar (\partial / \partial x)$$

Use this formula to write an expression for the part of the Hamiltonian that corresponds to the kinetic energy of a particle moving in the x direction. Show each step you take. (As you may recall, the Hamiltonian is the operator H which acts on the wavefunction to give the total energy times the wavefunction or; $H\psi = E\psi$). (4 points)

$KE = \frac{1}{2} m v^2$ $m v = p$ $KE = \frac{1}{2} p v$ $v = \frac{d}{dt}$ $\hbar = \frac{h}{2\pi}$
 $H = \frac{-i \hbar v}{2} \frac{\partial}{\partial x}$ ~~$v = \frac{d}{dt}$~~ +2
 $= \frac{-i \hbar}{2} \frac{\partial^2}{\partial x^2}$ $\frac{-i \hbar}{4\pi} \frac{\partial^2}{\partial x^2} = \frac{\pi}{1}$ \odot v is actually $\frac{dx}{dt}$

2(b). Explain and show mathematically why the following expression,

$$\psi(x) = A \sin kx$$

is a possible solution to the Schrödinger equation for a particle moving in a 1-D box. The Schrödinger equation for this system is,

$$\partial^2 / \partial x^2 (\psi(x)) = - [(8 \pi^2 m E) / h^2] \psi(x) \quad (4 \text{ points})$$

$\frac{d}{dx}$
 $\frac{d^2}{dx^2}$
 $\psi(x) = A \sin(kx)$
 $\psi'(x) = kA \cos(kx)$
 $\psi''(x) = -k^2 A \sin(kx)$

it fulfills the equation because the second derivative of the function is itself $\sin(kx)$ +4

2(c). If the kinetic energy of an electron is known to lie between $1.59 \times 10^{-19} \text{ J}$ and $1.61 \times 10^{-19} \text{ J}$ what is the smallest distance within which it can be known to lie (4 points)

$\lambda = \frac{h}{mv}$ $KE = \frac{1}{2} m v^2$ $\frac{2 KE}{v} = m v$ $v_1 = \sqrt{\frac{2 KE}{m_e}} = 591000 \text{ m/s}$
 $\lambda_1 = \frac{h}{m_e v_1} = 1.23 \text{ nm}$
 $\lambda_2 = \frac{h}{m_e v_2} = 1.22 \text{ nm}$

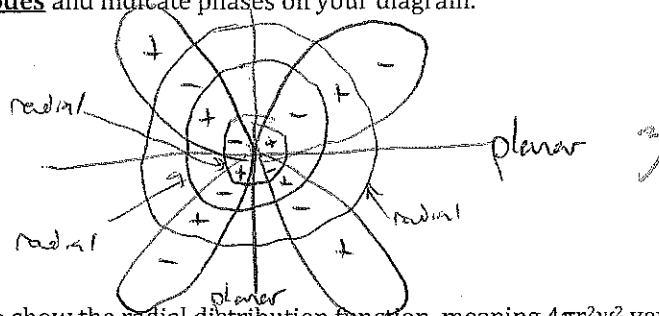
1.23
 -1.22
 $\boxed{.01 \text{ nm}}$

3(a). How many planar nodes and radial nodes are present in any 6d orbitals?

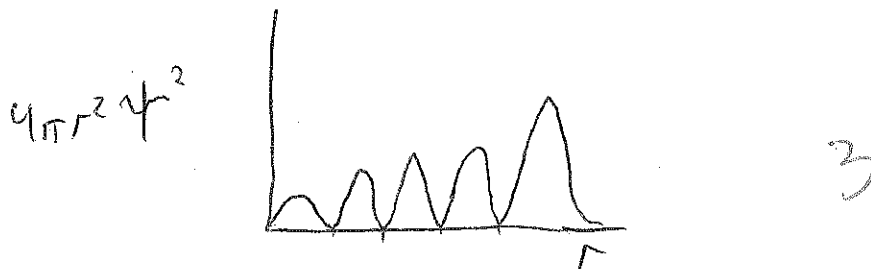
Planar.....2..... Radial3.....

0 1 2 3
s p d f
5 total
2 planar
(2)
3 radial
(3)

Draw **all the nodes** and indicate phases on your diagram.



3(b). Draw a graph to show the radial distribution function, meaning $4\pi r^2 \psi^2$ versus r , for a 5s orbital. (3)



3(c). What is the wavelength of a spectral line due to a transition between the $n = 3$ and $n = 7$ levels of an ion of O^{+7} ? (4)

$$E = -R_{\infty} (Z^2) \left(\frac{1}{9} - \frac{1}{49} \right) = \frac{hc}{\lambda} = 1.26 \times 10^{-17} \text{ J}$$

~~10 nm~~
15.7 nm

3(d) Explain why ionization energy shows a decrease between the following elements, (4)

(i) arsenic & selenium

Arsenic has a p orbital with 3 electrons one in each sublevel, adding another to get selenium puts two e^- into one sublevel increasing e^-e^- repulsion, decreasing energy

(ii) magnesium to aluminum

one e^- ~~more than~~ magnesium will create a new p shell and so elements in the same group as Al only have one e^- in outer shell which has lower ionization energy compared to the full s shell of Mg

9

12

Each part is worth 2 points.

- 4(a). Why is it strictly incorrect to write the configuration of manganese as $[\text{Ar}] 4s^2 3d^5$?

+ For manganese, the 3d orbital is lower energy than 4s.

- 4(b). What is the configuration of the Cu^+ ion?

+ $1s^2 2s^2 2p^6 3s^2 3p^6 3d^{10}$ (loses 4s)

- 4(c). What connection, if any, is there between the answer to 4b and part 4a?

+ For Cu^+ , it lost the 4s electron because it was higher energy than 3d, when you ionize you take away the outer most e^- which is why Mn should be $[\text{Ar}] 3d^5 4s^2$ to show it loses 4s e^- first.

- 4(d). Why is the configuration of potassium $[\text{Ar}] 4s^1$ rather than $[\text{Ar}] 3d^1$?

+ For potassium, the 4s orbital is lower energy than the 3d orbital so following Aufbau it goes into 4s first.

- 4(f). What is unique about the configuration of the palladium atom (Pd)?

+ It skips the 5s orbital completely. IDK!!!

- 4(g). Explain why the atom of yttrium (Y) can show greater stability by adopting a configuration of $[\text{Kr}] 4d^1 5s^2$ rather than $[\text{Kr}] 4d^3$?

+ The full 5s shell is less unstable compared to having 3 out of 10 in the d shell.