

27th February 2024

I have an ADA accommodation to do my assignment on paper. This document is a utilization of that accommodation. This assignment will utilize the makeup questions for Exam II provided by Dr. Moulder. Reasoning will be provided within the context of this class, and at times, within the context of concepts outside of the scope of this class. Each problem will appear on a new page.

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**Temperature Conversions**

$$^{\circ}F = \frac{9}{5}^{\circ}C + 32$$

$$^{\circ}C = \frac{5}{9} (^{\circ}F - 32)$$

$$K = 273.15 + ^{\circ}C$$

**Constants**

$$N = 6.0223 \times 10^{23}$$

$$c = 3.00 \text{ m s}^{-1}$$

$$h = 6.63 \times 10^{-34} \text{ J s}$$

**Energy and Electromagnetic Waves**

$$\nu = \frac{c}{\lambda}$$

$$E = \frac{hc}{\lambda}$$

$$E = h\nu \quad \text{Hz} = \text{per/second}$$

**Gas Law**

$$PV = nRT \quad \frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$

$$R = 8.21 \times 10^{-2} \text{ L atm mol}^{-1} \text{ K}^{-1}$$

$$\text{STP Gas Law} = 0^{\circ}\text{C}, 1 \text{ atm}$$

$$1 \text{ atm} = 760 \text{ Torr} = 760 \text{ mm Hg}$$

**Colligative Properties**

$$\Delta T_f = i K_f m$$

$$\Delta T_b = i K_b m$$

**Thermodynamics**

$$\Delta H_{rxn}^{\circ} = \sum n \Delta H_f^{\circ} - \sum m \Delta H_f^{\circ}$$

$$\Delta E = \Delta H - P \Delta V$$

$$q_1 = -q_2 \quad w = -P \Delta V \quad 1 \text{ L atm} = 101.3 \text{ J}$$

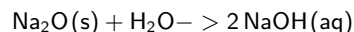
$$C = ms \quad q = ms \Delta t \quad E = q + w$$

**Insoluble Compounds****Soluble Compounds**

1. All salts of the alkali metals are soluble
2. All salts containing  $\text{NH}_4^+$ ,  $\text{NO}_3^-$ ,  $\text{ClO}_4^-$ ,  $\text{ClO}_3^-$  and  $\text{C}_2\text{H}_3\text{O}_2^-$  are soluble.
3. All chlorides, bromides, and iodides (salts containing  $\text{Cl}^-$ ,  $\text{Br}^-$ , or  $\text{I}^-$ ) are soluble except when combined with  $\text{Ag}^+$ ,  $\text{Pb}^{2+}$ , and  $\text{Hg}_2^{2+}$ .
4. All salts containing  $\text{SO}_4^{2-}$  are soluble except those of  $\text{Pb}^{2+}$ ,  $\text{Ca}^{2+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Ba}^{2+}$ , and  $\text{Hg}_2^{2+}$ .

1. All metal hydroxides (ionic compounds containing  $\text{OH}^-$ ) and all metal oxides (ionic compounds containing  $\text{O}^{2-}$ ) are insoluble except those of Group 1A and those of  $\text{Ca}^{2+}$ ,  $\text{Sr}^{2+}$ , and  $\text{Ba}^{2+}$ .

- When metal oxides do dissolve, they react with water to form hydroxides. The oxide ion,  $\text{O}^{2-}$ , does not exist in water. For example:



2. All salts containing  $\text{PO}_4^{3-}$ ,  $\text{CO}_3^{2-}$ ,  $\text{SO}_3^{2-}$ , and  $\text{S}^{2-}$  are insoluble except those of Group 1A and  $\text{NH}_4^+$

	Element	Oxidation Reaction
	Lithium	$\text{Li} \longrightarrow \text{Li}^+ + \text{e}^-$
	Potassium	$\text{K} \longrightarrow \text{K}^+ + \text{e}^-$
	Barium	$\text{Ba} \longrightarrow \text{Ba}^{2+} + 2 \text{e}^-$
	Calcium	$\text{Ca} \longrightarrow \text{Ca}^{2+} + 2 \text{e}^-$
	Sodium	$\text{Na} \longrightarrow \text{Na}^+ + \text{e}^-$
	Magnesium	$\text{Mg} \longrightarrow \text{Mg}^{2+} + 2 \text{e}^-$
	Aluminum	$\text{Al} \longrightarrow \text{Al}^{3+} + 3 \text{e}^-$
	Manganese	$\text{Mn} \longrightarrow \text{Mn}^{2+} + 2 \text{e}^-$
	Zinc	$\text{Zn} \longrightarrow \text{Zn}^{2+} + 2 \text{e}^-$
	Chromium	$\text{Cr} \longrightarrow \text{Cr}^{3+} + 3 \text{e}^-$
	Iron	$\text{Fe} \longrightarrow \text{Fe}^{2+} + 2 \text{e}^-$
	Cadmium	$\text{Cd} \longrightarrow \text{Cd}^{2+} + 2 \text{e}^-$
	Cobalt	$\text{Co} \longrightarrow \text{Co}^{2+} + 2 \text{e}^-$
	Nickel	$\text{Ni} \longrightarrow \text{Ni}^{2+} + 2 \text{e}^-$
	Tin	$\text{Sn} \longrightarrow \text{Sn}^{2+} + 2 \text{e}^-$
	Lead	$\text{Pb} \longrightarrow \text{Pb}^{2+} + 2 \text{e}^-$
	Hydrogen	$\text{H}_2 \longrightarrow 2 \text{H}^+ + 2 \text{e}^-$
	Copper	$\text{Cu} \longrightarrow \text{Cu}^{2+} + 2 \text{e}^-$
	Silver	$\text{Ag} \longrightarrow \text{Ag}^+ + \text{e}^-$
	Mercury	$\text{Hg} \longrightarrow \text{Hg}^{2+} + 2 \text{e}^-$
	Platinum	$\text{Pt} \longrightarrow \text{Pt}^{2+} + 2 \text{e}^-$
	Gold	$\text{Au} \longrightarrow \text{Au}^+ + \text{e}^-$

## Calculate the Velocity of Electrons with de Broglie Wavelengths of 285 nm.

The equation

$$\lambda = \frac{h}{mv}$$

is accurate for non-relativistic velocities. For relativistic velocities, the equation

$$\frac{1}{\lambda_B} = \frac{mv}{h\sqrt{1 - \frac{v^2}{c^2}}}$$

is more accurate. I will determine the velocity of electrons with de Broglie wavelengths of 285 nm using both of these equations.

### Calculating the de Broglie Wavelength with the Non-Relativistic Equation

For the non-relativistic equation, we will first consider that the mass of an electron is given to be  $9.1094 \times 10^{-31}$  kg. The non-relativistic equation:

$$\lambda = \frac{h}{mv}$$

can be rewritten:

$$h = mv\lambda.$$

To determine the velocity, we can rearrange the equation to give the velocity:

$$v = \frac{h}{m\lambda}.$$

Substituting the variables with known values, with the knowledge that  $1 \text{ J} = 1 \text{ kg m}^2 \text{ s}^{-2}$ , gives:

$$v = \frac{6.63 \times 10^{-34} \text{ kg m}^2 \text{ s}^{-2} \text{ s}}{9.1094 \times 10^{-31} \text{ kg} \times 2.85 \times 10^{-7} \text{ m}}.$$

This gives:

$$v \text{ m s}^{-1} = \frac{6.63 \times 10^{-34} \text{ kg m}^2 \text{ s}^{-1}}{2.596179 \times 10^{-37} \text{ kg m}}$$

which provides the value:

$$v = 2.553753035 \times 10^3 \text{ m s}^{-1} = 2.553 \times 10^3 \text{ m s}^{-1}$$

This value is closest to multiple choice option **D**:  $2.552 \times 10^3 \text{ m s}^{-1}$ .

### Calculating the de Broglie wavelength using the Proper Relativistic Equation

The de Broglie wavelength of a particle is a wavelength that determines the probability density of the particle being found at a specific point  $q$  in its configuration space. This is important in quantum mechanics. In the context of General Chemistry I, the de Broglie wavelength is defined in our textbook only as a characteristic of particles and other bodies. The relativistic definition of the de Broglie wavelength is:

$$\lambda_B = \frac{h}{p},$$

where  $\lambda_B$  represents the de Broglie wavelength of the particle,  $h$  represents the Planck constant, which has the value  $6.626\,070\,15 \times 10^{-34}$  Js, and  $p$  represents the relativistic momentum of the particle, defined as

$$p = \frac{mv}{\sqrt{1 - \frac{v^2}{c^2}}},$$

with  $m$  representing the mass of the particle,  $v$  representing the particle's velocity, and  $c$  representing the speed of light in a vacuum. The invariant mass of the electron is approximately  $9.109\,383\,701\,5(28) \times 10^{-31}$  kg according to the NIST. The speed of light in a vacuum is defined as  $299\,792\,458$  m s<sup>-1</sup>.

Combining these equations gives the formula:

$$\lambda_B = \frac{h}{\frac{mv}{\sqrt{1 - \frac{v^2}{c^2}}}}$$

which simplifies to

$$\lambda_B = \frac{h\sqrt{1 - \frac{v^2}{c^2}}}{mv}.$$

This can be rearranged to

$$\lambda_B mv = h\sqrt{1 - \frac{v^2}{c^2}}.$$

root -? The following steps will solve the equation for velocity:

$$\frac{\lambda_B mv}{h} = \sqrt{1 - \frac{v^2}{c^2}} \Rightarrow \left(\frac{\lambda_B mv}{h}\right)^2 = \left(\sqrt{1 - \frac{v^2}{c^2}}\right)^2 \Rightarrow \frac{\lambda_B^2 m^2 v^2}{h^2} = 1 - \frac{v^2}{c^2} \Rightarrow$$

$$1 = \frac{\lambda_B^2 m^2 v^2}{h^2} + \frac{v^2}{c^2} \Rightarrow 1 = \frac{\lambda_B^2 m^2 v^2 c^2 + v^2 h^2}{h^2 c^2} \Rightarrow h^2 c^2 = \lambda_B^2 m^2 v^2 c^2 + v^2 h^2 \Rightarrow$$

$$h^2 c^2 = v^2 (\lambda_B^2 m^2 c^2 + h^2) \Rightarrow v^2 = \frac{h^2 c^2}{\lambda_B^2 m^2 c^2 + h^2} \Rightarrow v = \sqrt{\frac{h^2 c^2}{\lambda_B^2 m^2 c^2 + h^2}}$$

$$v = \sqrt{\frac{h^2}{\lambda_B^2 m^2}}$$

And thus, we have arrived at the relativistic equation for the de Broglie wavelength solved for velocity. Using the values we defined earlier, we find the velocity to be:

$$v^2 = \frac{(6.626\,070\,15 \times 10^{-34} \text{ Js})^2}{(285 \times 10^{-9} \text{ m})^2 \times (9.109\,383\,701\,5 \times 10^{-31} \text{ kg})^2}$$

$$v^2 = \frac{4.390\,480\,563 \times 10^{-67} \text{ kg}^2 \text{ m}^4 \text{ s}^{-2}}{6.740\,121\,281 \times 10^{-74} \text{ kg}^2 \text{ m}^2}$$

$$v^2 = 6.513\,948\,904 \times 10^6 \text{ m}^2 \text{ s}^{-2}$$

$$v = 2.552\,243\,896 \times 10^3 \text{ m s}^{-1},$$

which, upon applying significant figures, would come to

$$v = 2.552 \times 10^3 \text{ m s}^{-1}$$

This is exactly the multiple choice option **D**.

### **Solution**

Both the non-relativistic approximation and the true relativistic equation give the answer

$$\mathbf{D.} \quad 2.552 \times 10^3 \text{ m s}^{-1}.$$

## **What Element is Theoretically the Smallest of all on the Periodic Table?**

The smallest theoretical element is Helium. This follows the periodic trends of atomic radius. As an element's group grows, its radius shrinks as there are more electrons in its shell, and the radius grows as the period increases, as there are more electron shells. Since Helium has the most electrons and the least electron shells, Helium is the smallest element.

**What wavelength would the hydrogen Rydberg line give for the electronic transition  $n_1 = 2$  and  $n_2 = 5$ ?**

The Rydberg formula models the emission and absorption wavelengths for photons in jumps in electron energy levels. It is defined for hydrogen as follows:

$$\frac{1}{\lambda_{vac}} = R_H \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right)$$

where  $R_H$  is defined in the handout as  $1.097 \times 10^7 \text{ m}^{-1}$ . This gives us the equation

$$\frac{1}{\lambda_{vac}} = (1.097 \times 10^7 \text{ m}^{-1}) \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right)$$

The problem is asking for the Rydberg line for  $n_1 = 2$  and  $n_2 = 5$ . Plugging these into the formula gives us

$$\frac{1}{\lambda_{vac}} = (1.097 \times 10^7 \text{ m}^{-1}) \left( \frac{1}{4} - \frac{1}{25} \right)$$

Fraction Subtraction rules tell us that we need to find a common denominator.

$$\frac{1}{\lambda_{vac}} = R_H \left( \frac{25}{100} - \frac{4}{100} \right) = R_H \left( \frac{21}{100} \right)$$

From this, we can see that

$$\lambda_{vac} = \frac{100}{21 \times 1.097 \times 10^7 \text{ m}^{-1}}$$

Which gives us

$$\lambda_{vac} = 4.341 \times 10^{-7} \text{ m}$$

which is approximately equal to 434.0 nm.

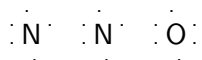
Thus, the wavelength of the hydrogen Rydberg line for the electronic transition from 2 to 5 would be 434.0 nm, or multiple choice option **C**.

## Which is the most correct Lewis structure for N<sub>2</sub>O

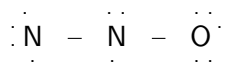
Both Nitrogen and Oxygen in N<sub>2</sub>O can be the central atom, so we determine the central atom by determining the least electronegative atom and placing it in the middle. N has electronegativity 3.04, while O has electronegativity 3.44. Thus, Nitrogen will be the central atom.

There are five valence electrons in each nitrogen atom, and six in the oxygen atom. Thus, there are sixteen total valence electrons in N<sub>2</sub>O.

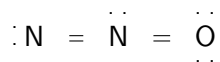
Next, we must draw the skeleton structure for N<sub>2</sub>O.



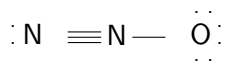
Now, we connect each atom with the central atom with a single bond.



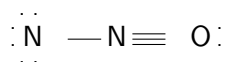
Now, we must create new bonds in the central atom to create double and triple bonds.



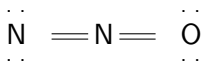
The N atom on the end does not follow the octet rule. Thus, we must move around bonds to make it valid.



This works, but there are resonance structures:



and



Now, we must compute the formal charges of each of the above resonance structures. Formal Charge can be calculated via the following

$$Q_{\text{Formal}} = V_{e^-} - LP_{e^-} - \frac{B_{e^-}}{2}$$

For the first structure, N $\equiv$ N—O, the Formal Charge is computed:

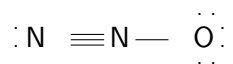
$$N_{\text{edge}} : 5 - 2 - \frac{6}{2} = 0$$

$$N_{\text{central}} : 5 - 0 - \frac{8}{2} = +1$$



$$\mathbf{O} : 6 - 6 - \frac{2}{2} = -1$$

Thus the formal charge of the first resonance structure,  $\text{N} \equiv \text{N} - \text{O}$ , is  $0 + 1 - 1 = 0$ . This has the negative charge on the Oxygen and the positive charge on the Nitrogen. The Oxygen is the more electronegative of the two, so this is the most correct structure for  $\text{N}_2\text{O}$ :



This version is close to multiple choice option **D**.

## What is the Frequency of Light that has a wavelength of 185 nm?

The frequency of a photon can be derived in a classical sense through

$$\lambda = \frac{c}{\nu}$$

as such

$$\nu = \frac{c}{\lambda}$$

We can input the necessary values for  $c$  and  $\lambda$ .

$$\nu = \frac{3.00 \times 10^8 \text{ m s}^{-1}}{185 \times 10^{-9} \text{ m}}$$

this gives us the value

$$\nu = 1.62 \times 10^{15} \text{ s}^{-1}$$

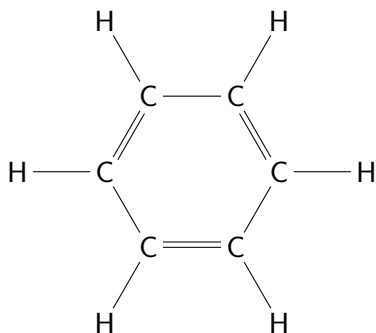
This is the value of the multiple choice option **E**.

**Which element is most likely to present with the first five ionization energies specified below? (in  $\text{kJ mol}^{-1}$ )**

589.8	1145	4912	6491	8153
-------	------	------	------	------

These five ionization energies are increasing in order. Since the third ionization energy is eemphibits a great increase, and it is greatly difficult to remove an electron from a full octet, one can assume that the atom has two valence electrons. Looking at the provided options, the only group 2 element in Calcium. Thus, one can assume the answer is Calcium, multiple choice option **B**.

**What is the hybridization of all the carbon atoms in benzene ( $\text{C}_6\text{H}_6$ )?**



The carbon atoms in benzene each have three electron regions, and thus each has the orbital hybridization  $sp^2$ .

I'm going to be honest. I just can't grasp the mechanism by which electron orbital hybridization occurs, because I do not understand the underlying mathematics of it. I have rote memorized the 'rule' of association between electron regions and orbital hybridization, but I want to understand it on a deeper level.

### **Solution**

**E.**  $sp^2$

## Based on their electronic configurations, which of the following elements or ions is paramagnetic in a vapor phase?

Paramagnetism is a weak attraction to magnetic fields because of the spin of electrons creating magnetic moments.

### Sulfur

Sulfur has the following valence electron spin configuration:

↑↓	↑↓	↑	↑
3s	3p		

Sulfur can have magnetic properties. Its electron spins, and thus electron magnetic moments, are unopposed and create an atomic magnetic momentum.

### Zinc<sup>2+</sup>

Zinc<sup>2+</sup> has the following valence electron spin configuration:

□	↑↓	↑↓	↑↓	↑↓	↑↓	□	□	□
4s	3d					4p		

Zinc<sup>2+</sup> cannot have magnetic properties. Its electron's spins, and thus magnetic moments, are opposed and equalize to no atomic magnetic moment.

### Krypton

Krypton has the following valence electron spin configuration:

↑↓	↑↓	↑↓	↑↓	↑↓	↑↓	↑↓	↑↓	↑↓
4s	3d					4p		

Krypton cannot have magnetic properties. Its electron's spins, and thus magnetic moments, are opposed and equalize to no atomic magnetic moment.

### Tin<sup>2+</sup>

Tin<sup>2+</sup> has the following valence electron spin configuration:

↑↓	↑↓	↑↓	↑↓	↑↓	↑↓	□	□	□
5s	4d					5p		

Tin<sup>2+</sup> cannot have magnetic properties. Its electron's spins, and thus magnetic moments, are opposed and equalize to no atomic magnetic moment.

### Oxygen<sup>2-</sup>

Oxygen<sup>2-</sup> has the following valence electron spin configuration:

↑↓	↑↓	↑↓	↑↓
2s	2p		

Oxygen<sup>2-</sup> cannot have magnetic properties. Its electron's spins, and thus magnetic moments, are opposed and equalize to no atomic magnetic moment.

### Solution

Sulfur is the only atom with a atomic magnetic moment. Thus, Sulfur, option **A.** is the correct answer.

**What is the maximum number of electrons that can have the following set of quantum numbers:  $n = 4$ ,  $l = 3$ ,  $m_l = 3$ ,  $m_s = -\frac{1}{2}$**

**ONE!!** Basic physics concepts dictate that no two electrons can inhabit the same quantum state. In non-relativistic quantum mechanics, there is no proof for the Pauli Exclusion Principle, though I believe one arises in relativistic quantum mechanics. I do not know that proof. However, the most basic physical principles dictate that only one electron can inhabit the same quantum state.

**Order the following series of isoelectronic ions ( $\text{Mg}^{2+}$ ,  $\text{N}^{3-}$ ,  $\text{F}^-$ ,  $\text{Si}^{4+}$ ) from largest to smallest ionic radii.**

When ionization of an atom occurs, an atom that loses an electron is called a cation. Likewise, an atom that gains an electron is called an anion. Because a cation loses an electron, its atomic radius is necessarily smaller than its parent atom. Likewise, anions have greater atomic radii than their parent atom. Because of this, one can assume that isoelectronic ions, or ions with the same electronic configuration, one can order isoelectronic ions in order of ionic radii according to their charges.

Thus, the order that ensures ionic radii are ordered from largest to smallest is  $\text{N}^{3-} > \text{F}^- > \text{Mg}^{2+} > \text{Si}^{4+}$ .

### **Solution**

The order that ensures ionic radii are ordered from largest to smallest is  $\text{N}^{3-} > \text{F}^- > \text{Mg}^{2+} > \text{Si}^{4+}$ .

This is equivalent to option **E**.