Dr. Zellmer
Time: 7 PM Sun.
40 min

Chemistry 1250
Spring Semester 2022
Quiz V

Name $\qquad$
KEY
Lab TA/time

T, R
February 20, 2022

1. (5 pts) For the Hydrogen atom, calculate the wavelength (in nm ) of a photon for a transition from the $7^{\text {st }}$ shell to the $2^{\text {nd }}$ shell? Is this photon being emitted or absorbed by the atom? Is this in the visible portion of the electromagnetic spectrum (and how do you know)? (Show work and explain your answers.)

Use the Rydberg Eqn.

$$
--\frac{1}{\lambda}=R_{H}\left(\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right)
$$

This is technically for the wavelength of a photon emitted $\left(\mathrm{n}_{\mathrm{f}}<\mathrm{n}_{\mathrm{i}}\right)$ so it comes out positive. For an absorption $\left(n_{f}>n_{i}\right)$ it would come out negative but you ignore the negative sign since the energy of a photon can't be negative (take the absolute value) and just realize it's an absorption.

$$
\begin{aligned}
\frac{1}{\lambda} & =\left(1.097 \times 10^{7} \mathrm{~m}^{-1}\right)\left|\left(\frac{1}{2^{2}}-\frac{1}{7^{2}}\right)\right|=2.51 \underline{\mathbf{8}} 62 \times 10^{6} \mathrm{~m}^{-1} \\
\lambda & =3.97 \underline{0} 42 \times 10^{-7} \mathrm{~m}=397 \mathrm{~nm}
\end{aligned}
$$

The photon is being emitted since the transition is from $n=7$ to $n=2$.

This would NOT be in the visible region of the spectrum. The visible region is $400-750 \mathrm{~nm}$. This would be in the Ultraviolet (UV) region.
2. (3 pts) Which of the following electron transitions requires the smallest energy to be absorbed by the hydrogen atom?
a) from $\mathrm{n}=1$ to $\mathrm{n}=2$
b) from $n=2$ to $n=4$
c) from $\mathrm{n}=2$ to $\mathrm{n}=6$
d) from $\mathrm{n}=3$ to $\mathrm{n}=6$
e) * from $n=4$ to $n=6$

Remember, the energy diagram. The further the energy levels get from the nucleus the closer together they get in terms of energy. There's a big gap between $n=1$ and $n=2$, the gap between $n=2$ and $n=3$ is smaller, and so on and so forth.

From $n=4$ to $n=6$ would be the smallest gap, smallest energy needed to be absorbed.
3. ( 5 pts ) Calculate the de Broglie wavelength for a Ne molecule moving at $0.850 \mathrm{~m} / \mathrm{s}$ ? If this wavelength corresponded to a photon of light would we be able to see it in the visible portion of the electromagnetic spectrum (and how do you know)?

$$
\lambda=\frac{\mathrm{h}}{\mathrm{mv}} \quad \text { where } \mathrm{h}=\text { Planck's constant }\left(6.626 \times 10^{-34} \mathrm{~J} \bullet \mathrm{~s}\right), \mathrm{m}=\text { mass and } \mathrm{v}=\text { velocity }
$$

Need the mass of 1 Ne arom (20.179 amu) in $\mathrm{kg}, \quad\left(1 \mathrm{~J}=1 \mathrm{~kg} \cdot \mathrm{~m}^{2} \cdot \mathrm{~s}^{-2}\right)$

$$
\begin{aligned}
& ? \mathrm{~kg}=20.179 \mathrm{amu} \times \frac{1.66 \times 10^{-24} \mathrm{~g} \quad 1 \mathrm{~kg}}{1 \mathrm{amu}}=3.3497 \times 10^{-26} \mathrm{~kg} \\
& \lambda=\frac{\mathrm{h}}{\mathrm{mv}}=\frac{\left(6.626 \times 10^{-34} \mathrm{~J} \cdot \mathrm{~s}\right)\left[\left(1 \mathrm{~kg} \cdot \mathrm{~m}^{2} \cdot \mathrm{~s}^{-2}\right) / 1 \mathrm{~J}\right]}{\left(3.3 \underline{4} 97 \times 10^{-26} \mathrm{~kg}\right)(0.850 \mathrm{~m} / \mathrm{s})} \\
& \lambda=2.3 \underline{\mathbf{2}} 71 \times 10^{-8} \mathrm{~m}=23 . \underline{2} 7 \mathrm{~nm}=23.3 \mathrm{~nm}
\end{aligned}
$$

You would NOT be able to see this in the visible region of the spectrum. The visible region is 400-750 nm. This would be in the UV region (borderline with X-ray region).

## 4. The uncertainty principle states that

a) matter and energy are really the same thing
b) it is impossible to know how many electrons there are in an atom
c) there can only be one unknown digit in a number
d) it is impossible to know what orbitals the electrons occupy.
e)* it is impossible to know exactly both the position and momentum of an electron.

The Heisenberg Uncertainty Principle states it is impossible to determine exactly both the position (location) and momentum (velocity) of a particle. It applies to all particles but is only important when the masses are extremely small (subatomic particles). It can be stated in the following mathematical way:

$$
\begin{aligned}
& \Delta \mathrm{x} \cdot \Delta(\mathrm{~m} \bullet \mathrm{v})>-\frac{\mathrm{h}}{4 \pi} \\
& \Delta \mathrm{x}=\text { uncertainty in position } \\
& \Delta(\mathrm{mv})=\text { uncertainty in momentum }(\mathrm{p}=\mathrm{mv}) \\
& \mathrm{m}=\text { mass of the particle } \\
& \mathrm{v}=\text { velocity of particle } \\
& \mathrm{h}=\text { Planck's constant }=6.626 \times 10^{-34} \mathrm{~J} \bullet \mathrm{~S} \quad\left(1 \mathrm{~J}=1 \mathrm{~kg} \cdot \mathrm{~m}^{2} / \mathrm{s}^{2}\right)
\end{aligned}
$$

This means if either the position or momentum are known exactly the other has a minimum uncertainty greater than $\mathrm{h} / 4 \pi$.
5. (2 pts) Which of the following sets of quantum numbers are not allowed for an electron in an atom?

$$
\begin{array}{lcccc}
\text { 1)* } & \mathrm{n}=2 & \ell=2 & \mathrm{~m}_{\ell}=-1 & \mathrm{~m}_{\mathrm{s}}=+1 / 2 \\
\text { 2)* } & \mathrm{n}=3 & \ell=1 & \mathrm{~m}_{\ell}=+2 & \mathrm{~m}_{\mathrm{s}}=+1 \\
\text { 3) } & \mathrm{n}=4 & \ell=3 & \mathrm{~m}_{\ell}=-2 & \mathrm{~m}_{\mathrm{s}}=-1 / 2 \\
\text { 4) } \mathrm{n}=8 & \ell=6 & \mathrm{~m}_{\ell}=0 & \mathrm{~m}_{\mathrm{s}}=+1 / 2
\end{array}
$$

See \#11 above for a description of the quantum numbers.
$\mathrm{n}=1,2,3, \ldots, \infty \quad \ell=0,1,2, \ldots,(\mathrm{n}-1) \quad \mathrm{m}_{\ell}=-\ell, \ldots, 0, \ldots,+\ell \quad \mathrm{m}_{\mathrm{s}}=+1 / 2$ and $-1 / 2$

## Correct q.n.:

For (3) when $n=4$ and $\ell=3$ the possible values of $m_{\ell}$ are $m_{\ell}=-3,-2,-1,0,1,2,3$ and $m_{s}=-1 / 2$ ok
For (4) when $n=8$ and $\ell=6$ the possible values of $m_{\ell}$ are $m_{\ell}=-6, \ldots, 0, \ldots, 6$ and $m_{s}=+1 / 2 \mathrm{ok}$ Incorrect q.n.:

For (1) when $\mathrm{n}=2$ and $\ell=0,1$ ( $\ell$ can't be 2 when $\mathrm{n}=2$ )
For (2) when $n=3$ and $\ell=1$ can't have $m_{\ell}=+2$ or $m_{s}=1\left(m_{s}\right.$ can be only $+1 / 2$ and $\left.-1 / 2\right)$
6. Select an answer which includes all of the CORRECT statements given below.

1) There are nine $f$ orbitals in an f subshell.
2) A 10p subshell can have a maximum of $\mathbf{6}$ electrons.
3) Each $\mathbf{p}$ orbital within a subshell consists of four lobes along the axes.
4) There are 11 subshells in the $11^{\text {th }}$ shell.
5) For the ground state of an atom, within a shell, all the orbitals of a subshell are filled prior to putting electrons in the next subshell.

The correct statements are $2,4 \& 5$
2) A p subshell can have a maximum of $\mathbf{6}$ electrons. (True for all $p$ subshells, no matter what shell).
4) There are 11 subshells in the $11^{\text {th }}$ shell. (\# subshells in a shell $=$ shell \#, $\left.\mathbf{n}\right)$. There are $11^{2}(121)$ orbitals.
5) In the ground state of an atom, all orbitals within a subshell are filled prior to putting electrons in the next subshell.

The corrected answers for $1 \& 3$ would be:

1) There are seven f orbitals in an $f$ subshell.
2) Each p orbital within a subshell consists of two lobes. (figure eight, $\infty$; dumbbell)

The following table shows values of q.n. $\ell$, corresponding letters, \# orb. and max. \# e- in subshell

| $\ell=$ | 0 | 1 | 2 | 3 | 4 | $5 \ldots$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| subshell letter | s | p | d | f | g | h. . |
| \# orbitals in subshell | 1 | 3 | 5 | 7 | 9 | $11 \ldots$ |
| max. \# $\mathrm{e}^{-}$in subshell | 2 | 6 | 10 | 14 | 18 | $22 \ldots$ |

7. The electron configuration for an excited state of Gd is $[\mathrm{Xe}] 6 \mathrm{~s}^{1} 4 f^{7} 5 \mathrm{~d}^{2}$. How many unpaired electrons are in this excited atom? (Be careful!)
$[\mathrm{Xe}] \mathbf{6} \mathbf{s}^{\mathbf{1}} \mathbf{4 f}^{\mathbf{7}} \mathbf{5 d} \mathbf{d}^{\mathbf{2}} \quad$ or $\quad[\mathrm{Xe}] \mathbf{4} f^{\mathbf{7}} \mathbf{5 d}^{\mathbf{2}} \mathbf{6 s}^{\mathbf{1}}$
If the \# $\mathrm{e}-$ is $\preceq \# \mathrm{e}-$ in a half-filled subshell, then;
the number of unpaired $\mathrm{e}-=$ the number of $\mathrm{e}-\mathrm{in}$ the subshell.
If the \# $\mathrm{e}-$ is $>$ half filled (or filled), then;
\# of unpaired $\mathrm{e}-=$ (the maximum \# of $\mathrm{e}-$ in the subshell) - (the \# of $\mathrm{e}-$ present in the subshell).
There are 7 unpaired e- in the $4 \mathrm{f}, 2$ unpaired e-in the $5 \mathrm{~d} \& 1$ unpaired $\mathrm{e}-$ in the 6 s .
This gives a total of $\mathbf{1 0}$ unpaired $\mathrm{e}-$.

8. Which of the following electron configurations corresponds to an element in the same group as an element with a $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{6} 5 s^{2} 4 d^{10} 5 p^{2}$ electron configuration?
a) $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{4}$
b) $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{4}$
c) $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{6} 5 s^{2} 4 d^{2}$
d) $1 \mathrm{~s}^{2} 2 \mathrm{~s}^{2} 2 \mathrm{p}^{3}$
e)* $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{2}$

Both $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{2}$ and $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{6} 5 s^{2} 4 d^{10} 5 p^{2}$ are in Group 4A (IVA). (There are $4 \mathrm{e}-$ in the valence shell, the one with the highest $n$ value, $4 \mathrm{~s}^{2} 4 \mathrm{p}^{2}$ and $5 \mathrm{~s}^{2} 5 \mathrm{p}^{2}$.)
9. (4 pts) Fill in the blanks with the correct answer to each of the following.
(a) Which has the smaller ionization energy, ${ }_{32} \mathrm{Ge}$ or ${ }_{15} \mathrm{P}$ ?

(b) Which is largest (size): ${ }_{35} \mathrm{Br}^{-},{ }_{34} \mathrm{Se}^{2-}$ or ${ }_{33} \mathrm{As}^{3-}$ ?
has $36 \mathrm{e}^{-}$(like other two) but ${ }_{33} \mathrm{As}^{3-} 33$ protons (fewer protons) anions larger than cations in isoelectronic series (same \# e ${ }^{-}$); ${ }_{33} \mathrm{As}^{3-}$ has higher (-) charge (fewer protons for the same \# of electrons - things spread out)
(c) Which reacts more readily with chlorine, ${ }_{37} \mathrm{Rb}$ or ${ }_{55} \mathrm{Cs}$ ?
 (i.e. lose an $\mathrm{e}^{-}$more easily)

These metals would lose $\mathrm{e}-$ to Cl (more metallic react faster)
(d) Which has the more negative electron affinity, ${ }_{34} \mathrm{Se}$ or ${ }_{17} \mathrm{Cl}$ ?(several exceptions)

10. (3 pts) Explain how the effective nuclear charge, $\mathrm{Z}_{\mathrm{eff}}$, changes going across a row from left to right in the periodic table and why?

Going left to right across a row the $\mathrm{Z}_{\text {eff }}$ increases (rather rapidly). We are using the following for $\mathrm{Z}_{\text {eff }}$,

$$
\mathrm{Z}_{\mathrm{eff}}=\mathrm{Z}-\mathrm{S} \quad \mathrm{Z}=\# \text { protons } \quad \mathrm{S}=\text { Shielding factor }
$$

There are many ways to determine $S$. We're using the simplest one, $S$ equals the number of core electrons. Going across a row the number of core electrons remains constant. However, the \# protons increases by 1 each time you move one column to the right. Since $S$ is constant and $Z$ inc. by 1 when moving over 1 column the Zeff increases by 1 each time you move one column to the right. For example, in row 2 the core electrons are the two electrons in the 1 s (He core). As you move over one column to the right $Z$ inc. by 1 but the $\#$ of core electrons, $S$, stays constant at 2 and thus the $\mathrm{Z}_{\text {eff }}$ inc. by 1 . By the time you get to fluorine the $\mathrm{Z}_{\text {eff }}$ is +7 . This has big effects on the size and I.E.

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11. (3 pts) Explain why the electron affinity for Beryllium is positive while that for Boron is negative.

EA is the energy which occurs when an electron is added to a gaseous atom. Look at the $\mathrm{e}^{-}$ configurations for each atom.

B: $[\mathrm{He}] 2 \mathrm{~s}^{2} 2 \mathrm{p}^{1} \quad \mathrm{Be}:[\mathrm{He}] 2 \mathrm{~s}^{2}$
In Be the 2 s subshell is filled. A filled subshell is pretty stable. The added $\mathrm{e}^{-}$must be put in a previously empty p subshell that is higher in energy. Also, the $\mathrm{e}-$ which would go into the 2 p is shielded a little by the existing 2 s electrons. All this requires energy. In B the extra $\mathrm{e}^{-}$goes in an empty 2 p orbital in the 2 p subshell which already has $1 \mathrm{e}^{-}$to begin with and there is no extra shielding due to the existing $\mathrm{e}^{-}$in the 2 p subshell.
12. (2 pts) Write the equation corresponding to the first electron affinity of Oxygen.

Electron affinity: energy change which occurs when an electron is added to a gaseous neutral atom or anion. The first EA is for the first electron being added. For most neutral atoms energy is released (negative EA) when an electron is added. There are a few cases when the energy change is positive (such as the noble gases, $\mathrm{Be}, \mathrm{Mg}$ and N , the reasons for this were discussed in lecture and the textbook).

$$
\mathrm{O}(\mathrm{~g})+\mathrm{e}^{-} \rightarrow \mathrm{O}^{-}(\mathrm{g}) \quad \text { first E.A. (First electron being added.) }
$$

13. (2 pts) Write the equation corresponding to the second ionization energy of Ca .

Ionization: losing one or more electron from a gaseous atom or ion.
Ionization energy: energy required to remove an electron from a gaseous neutral atom or ion.

$$
\begin{array}{ll}
\mathrm{Ca}(\mathrm{~g}) \rightarrow \mathrm{Ca}^{+}(\mathrm{g})+\mathrm{e}^{-} & \text {first I.E. } \\
\mathrm{Ca}^{+}(\mathrm{g}) \rightarrow \mathrm{Ca}^{2+}(\mathrm{g})+\mathrm{e}^{-} & \text {second I.E. }
\end{array}
$$

14. (3 pts) Write the electron configuration for the following ion by starting with the electron configuration for the neutral atom and then the ion given. Show work.
$\mathrm{Fe}^{2+}$

Fe:
Shorthand config: $\quad[\mathrm{Ar}] 4 \mathrm{~s}^{2} 3 \mathrm{~d}^{6}$
Use noble gas from previous period to represent the core (inner) $e^{-}$.
or
[Ar] $3 \mathrm{~d}^{6} 4 \mathrm{~s}^{2} \quad$ is okay $<$ - shows val. shell orbitals together on "outside"
Orbital diagram: $[\mathrm{Ar}] \frac{1 \mathrm{l}}{4 \mathrm{~s}} \quad \frac{1 \mathrm{l}}{3 \mathrm{~d}} \frac{1}{3} \quad \frac{1}{-}$
In forming ions for transition metals electrons come out of the ns orbital before the (n-1)d orbitals. Remove the two electrons in the 4 s . For $\mathrm{Fe}^{3+}$ remove one electron from the 3d.
$\mathrm{Fe}^{2+} \quad[\mathrm{Ar}] 3 \mathrm{~d}^{6}$
Orbital diagram:

15. (3 pts) The electrons that are removed from ${ }_{48} \mathrm{Cd}$ to form the $\mathrm{Cd}^{2+}$ ion are from the $\qquad$ subshell and the electrons that are removed from ${ }_{83} \mathrm{Bi}$ to form the $\mathrm{Bi}^{2+}$ ion are from the $\qquad$ subshell.
a) * 5 s; 6 p
b) $5 \mathrm{~s} ; 5 \mathrm{~s}$
c) $4 \mathrm{~d} ; 4 \mathrm{f}$
d) $5 \mathrm{~d} ; 6 \mathrm{p}$
e) $4 \mathrm{~s} ; 5 \mathrm{~d}$

Need to write out the e-configuration for each atom and then look at which electrons will come out when it forms an ion. Remember, the valence $\mathrm{e}^{-}$come out first when forming ions.
$\mathrm{Cd} \quad[\mathrm{Kr}] 5 \mathrm{~s}^{2} 4 \mathrm{~d}^{10}$
$\mathrm{Cd}^{2+} \quad[\mathrm{Kr}] 4 \mathrm{~d}^{10} \quad$ electrons come out of the 5 s (valence shell) first
$\mathrm{Bi} \quad[\mathrm{Xe}] 6 \mathrm{~s}^{2} 4 \mathrm{f}^{14} 5 \mathrm{~d}^{10} 6 \mathrm{p}^{3}$
$\mathrm{Bi}^{2+} \quad[\mathrm{Xe}] 6 s^{2} 4 \mathrm{f}^{14} 5 \mathrm{~d}^{10} 6 \mathrm{p}^{1} \quad$ electrons come out of the 6 p (valence shell) first

