## CHEM1611 Worksheet 2 - Answers to Critical Thinking Questions

The worksheets are available in the tutorials and form an integral part of the learning outcomes and experience for this unit.

## Model 1: Atomic Orbitals

1. $s$ orbitals consist of a single lobe. $p$ orbitals consist of two large lobes. $d$ orbitals consist of four lobes. (The shape of $3 d_{\mathrm{z}} 2$ is a little different to that of the other $3 d$ orbitals. It consists of two large lobes with a ring around the centre.)
2. The orbital quantum number, $l$.
3. See table below.

| $\boldsymbol{n}$ | Possible $\boldsymbol{l}$ values | Orbital labels |
| :---: | :---: | :---: |
| 1 | 0 only | $1 s$ |
| 2 | 0 and 1 | $2 s$ and $2 p$ |
| 3 | 0,1 and 2 | $3 s, 3 p$ and $3 d$ |

4. See table below.
5. See table below.

| $\boldsymbol{l}$ | Possible $\boldsymbol{m}_{\boldsymbol{l}}$ values | Number of values of $\boldsymbol{m}_{\boldsymbol{l}}$ | Number of orbitals | Maximum number of <br> electrons |
| :---: | :---: | :---: | :---: | :---: |
| 0 | 0 | 1 | 1 | $1 \times 2=2$ |
| 1 | 1,0 and -1 | 3 | 3 | $3 \times 2=6$ |
| 2 | $2,1,0,-1$ and -2 | 5 | 5 | $5 \times 2=10$ |

## Model 2: Electronic Configurations of Atoms

1. 

(a) $1 s^{2} 2 s^{1}$ or $[\mathrm{He}] 2 s^{1}$
(b) $1 s^{2} 2 s^{2}$ or $[\mathrm{He}] 2 s^{2}$
(c) $1 s^{2} 2 s^{2} 2 p^{1}$ or $[\mathrm{He}] 2 s^{2} 2 p^{1}$
(d) $1 s^{2} 2 s^{2} 2 p^{2}$ or $[\mathrm{He}] 2 s^{2} 2 p^{2}$
(e) $1 s^{2} 2 s^{2} 2 p^{3}$ or [He] $2 s^{2} 2 p^{3}$
(f) $\quad 1 s^{2} 2 s^{2} 2 p^{4}$ or [He $2 s^{2} 2 p^{4}$
(g) $1 s^{2} 2 s^{2} 2 p^{5}$ or $[\mathrm{He}] 2 s^{2} 2 p^{5}$
(h) $\quad 1 s^{2} 2 s^{2} 2 p^{6}$ or $[\mathrm{He}] 2 s^{2} 2 p^{6}$
2. $\quad 1 s^{2} 2 s^{2} 2 p^{6} 3 s^{1}$ or $[\mathrm{Ne}] 3 s^{1}$. Both Na and Li have 1 electron in an $s$ orbital outside a noble gas configuration.
3. $2 s$ is lower in energy than $2 p$. ( $2 s$ is filled before $2 p$ is.).
4. The arrows refer to $m_{s}$ - the spin quantum number. According to the Pauli Principle, a maximum of one 'up' and one 'down' spin (arrow) can occupy each orbital (box).
5. Hund's Rule.
6. See below.

7. (i) It is quicker, (ii) it makes the valence electrons obvious and (ii) it shows the relationship between the valence configurations of elements.

The X atoms are the noble gases.
8. $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{6}$ or $[\mathrm{Ar}] 4 s^{2} 3 d^{6}$.
9. $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{3}$ or $[\mathrm{Ar}] 4 s^{2} 3 d^{10} 4 p^{3}$

## Model 2: Electronic Configurations of Atoms

1. The configurations of $\mathrm{Mg}^{2+}$ and $\mathrm{O}^{2-}$ are the same and match the preceding and proceeding noble gas respectively.

| Atom / Ion | Electron configuration | Representation - only valence electrons |  |
| :---: | :---: | :---: | :---: |
| O | $1 s^{2} 2 s^{2} 2 p^{4}$ or [He] $2 s^{2} 2 p^{4}$ | $\uparrow \downarrow$ | $\uparrow \downarrow$ ¢ $\uparrow$ |
| $\mathrm{O}^{2-}$ | $1 s^{2} 2 s^{2} 2 p^{6}$ or [He] $2 s^{2} 2 p^{6}$ | $\uparrow \downarrow$ | $\uparrow \downarrow \|$ |
| Mg | $\begin{gathered} 1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} \text { or } \\ {[\mathrm{Ne}] 3 s^{2}} \end{gathered}$ | $\uparrow \downarrow$ |  |
| $\mathrm{Mg}^{2+}$ | $\begin{gathered} 1 s^{2} 2 s^{2} 2 p^{6} \text { or } \\ {[\mathrm{Ne}]} \end{gathered}$ |  |  |

2. 

(a) $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6}$ or $[\mathrm{Ne}] 3 s^{2} 3 p^{6}$
(b) $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2}$ or $[\mathrm{Ne}]$
(c) It does not have one! $1 s^{0}$
(c) $1 s^{2}$

## Extension questions

1. As H is $1 s^{1}$ and He is $1 s^{2}$, a Periodic Table based purely on electronic configurations would have the elements in the $s$-block, above Li and Mg respectively.
The Periodic Table, however, was worked out from experimental observations of chemical and physical properties. He is an unreactive gas and for this reason, it is placed in the Nobel Gases. This can perhaps be justified from its electronic configuration as it has the maximum number of electrons possible for an element with $n=1$ just as Ne has the maximum number of electrons possible for an element with $n=2$.
H is more awkward! Chemically it behaves a little like the Group 1 elements - its commonest oxidation number is +1 , just like the Group 1 elements. However, compounds containing it with this oxidation number are very different to those of Group 1 elements: $\mathrm{LiOH}, \mathrm{NaOH}$ and KOH contain $\mathrm{M}^{+}$and $\mathrm{OH}^{-}$ions and are strongly basic. HOH is a covalent molecule and is not basic.
In its elemental form, hydrogen exists as a gas made up of $\mathrm{H}_{2}$ molecules whereas the Group 1 elements exist as metallic solids. (Note however that at very high pressure such as in the cores of planets, hydrogen probably is also metallic).
In forming the diatomic $\mathrm{H}_{2}$ molecule, hydrogen acts more like the halogens $\left(\mathrm{F}_{2}, \mathrm{Cl}_{2}, \mathrm{Br}_{2}\right.$ and $\mathrm{I}_{2}$ ). When it reacts with metals, it forms hydrides such as LiH which contain $\mathrm{H}^{-}$ions. This is akin to the halogens which also forms anions in compounds like LiF and NaCl .
There is no correct answer!
2. There are a number of trends:
(a) Ionisation energies decrease down each group as the electron to be removed is in an orbital with a higher $n$ quantum number and orbits further from the nucleus. As a result, the attraction to the nucleus is reduces.
(b) Ionisation energies generally increase across a period as the $n$ quantum number does not change but the positive charge of the nucleus increases. As a result, the attraction to the nucleus increases.
The combination of (a) and (b) means that the first element in each row ( $\mathrm{H}, \mathrm{Li}, \mathrm{Na}$ ) have the lowest ionisation energy in that row whilst the last element in each row ( $\mathrm{He}, \mathrm{Ne}$ and Ar ) have the highest ionisation energy in that row.
(c) Trend (b) is broken between the between Be and B and between Mg and Al . This is due to the change in the orbital of the electron which is being removed. When $\operatorname{Be}\left(2 s^{2}\right)$ is ionised, an $s$ electron is removed. When $\mathrm{B}\left(2 s^{2} 2 p^{1}\right)$ is ionised, a $p$ electron is removed. As $2 p$-orbitals have higher energies than $2 s$ orbitals, it is easier to ionise B than Be despite the nuclear charge being higher.
(d) Trend (b) is also broken between N and O and between P and S . This is due to the change in the spin of the electron being ionised. In $\mathrm{N}\left(2 s^{2} 2 p^{3}\right)$, the $p$-electron being ionised has the same spin as the other $p$-electrons. In $\mathrm{O},\left(2 s^{2} 2 p^{4}\right)$, the $p$-electron being ionised has the opposite spin as the other $p$-electrons. (Draw the electronic configurations using the box notation to see this.)
Electrons with the opposite spin repel each other more than electrons with the same spin. It is then easier to remove an electron from O .
