

2.1 THE NUCLEAR ATOM

- mass of an atom is in its nucleus, +ve nucleus
- nature of science → plum pudding model ⇒ gold leaf experiment (Rutherford)
 - subatomic particle theory was a paradigm shift in 1800s
- bohr model of an atom: a small solar system with electrons orbiting nucleus
neutrons needed to stabilize nuclei (+ve repel = fall apart)

a nucleus is composed of protons + neutrons [this is the mass number], (Z)

↳ electrons occupy space around the nucleus (outside)

· atomic number = number of protons. (A)

A \times n → charge

Z \times e.g. ${}^7_3\text{Li}$ has 3 protons and 4 neutrons, 3 electrons.

isotopes are atoms of the same element with different numbers of neutrons.

↳ different mass numbers. (diff physical properties - mp, bp, diff. rate)

relative atomic mass average mass of ALL isotopes of an element

compared to an atom of C-12

e.g. $(\% \times \text{mass}) + (\% \times \text{mass})$

100

or if you're trying to find indiv mass (100-x)

chemical properties of isotopes are the same as they have same electronic configuration and electrons in outer shell. [USE MASS SPECTROMETER + RAM]

a radioisotope has an unstable nucleus and undergoes radioactive decay

↳ emits gamma/alpha/beta particle

↳ used in medicine = diagnosis/treatment/research

e.g. tracers e.g. radiocarbon dating

↳ PET scans = 3D images of tracer concentration in the body (cancer)

e.g. Iodine-131 (Positron emission tomography) injected into blood, absorbed by good tissue, scanner detects photons of unabsorbed.

· RELATIVE ATOMIC MASS found via a mass spectrometer - determines abundance of isotope in an element to work out RAM

2.2 ELECTRON ARRANGEMENT

- electrons are arranged in **energy levels** of increasing energy and distance from the nucleus (first energy level, $n=1$, is closest to nucleus)

↳ it has the **lowest** energy

↳ can contain **max 2** electrons. ($n=2$, max 8, $n=3$, max 18)

- electrons in the outermost energy level = **valence electrons**

↳ they define the group number of the element.

evidence for energy levels come from the **VINE EMISSION SPECTRUM**

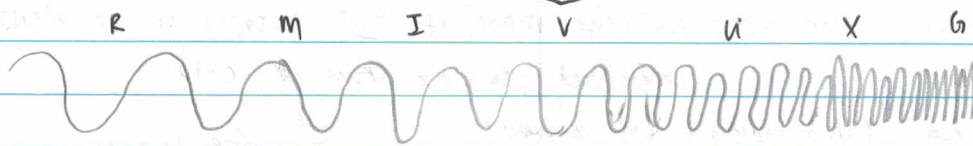
1. when electrons are excited = move from ground state to **HIGHER** energy level.

2. when it drops back into a lower shell = emit energy in form of light (photon)

↳ what could it release?

anything on the electromagnetic spectrum.

RO γ G B I V (colours)



low frequency

long wavelength

[energy \propto frequency
frequency \propto $1/\text{wavelength}$]

high frequency

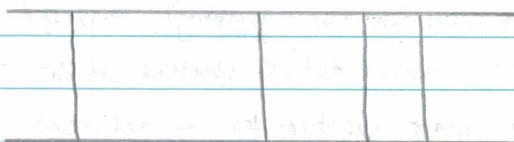
short wavelength

continuous spectrum: all possible wavelengths/frequencies

2 (a). when energy is supplied / released, the element will emit a spectrum of **CERTAIN** lines at **CERTAIN** wavelengths.

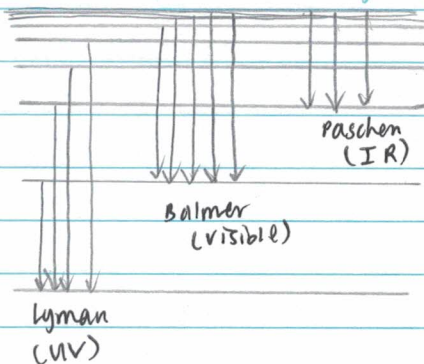
the **line spectrum** is a characteristic spectrum of discrete lines.

line emission spectrum of hydrogen



- this is evidence of electrons in discrete energy levels which **CONVERGE** at higher energies.

lower energy
emission spectrum of hydrogen



- the jump back to $n=1$ is the largest drop in energy. High energy, high frequency, UV.
- when electron jumps to $n=2$ level, this is seen in (VISIBLE) region
- each element has different energy level distances = diff. line spectrum.

2.2 ELECTRON ARRANGEMENT (CONT'D)

each energy level is n and has a number (1st is $n=1$ etc)

each energy level can hold $2n^2$ number of electrons.

within each energy level, there are subshells.
 there are FOUR types of subshells: s, p, d, f .
 each sub-shell is made of electron orbitals which hold 2 electrons.

this is where an electron is likely to be.

s subshells have lowest energy, then p , then d , then f .
 \downarrow \downarrow \downarrow
 1 orbital 3 orbitals 5 orbitals

In each energy level ...
 $1s$ ($n=1$) has 1 subshell (s)
 $(n=2)$ has 2 subshells (s, p)
 $n=3$ has 3 subshells (s, p, d)

each orbital has a different shape

$\hookrightarrow s =$ spherical (0)

$\hookrightarrow p_{x, y, z} =$ dumbbell (8) $x = \infty$ $y = \infty$ $z = \infty$

electron configuration

① aufbau principle

orbitals of lowest energy = filled first

② Hund's rule

each orbital must have ONE electron b4 pairing.

e.g. He = $1s^2$ $1 =$ energy level
 $s =$ subshell
 $2 =$ no. of e^-

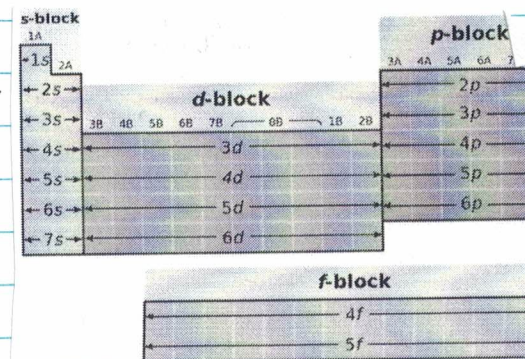
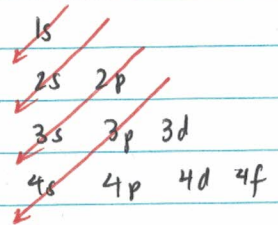
e.g. Li = $1s^2 2s^1$ or 'electrons in boxes' diagrams
 $N = 1s^2 2s^2 2p^3$

because $3d$ has more energy than $4s$, $3d$ AFTER $4s$.
 Ne = $1s^2 2s^2 2p^6 3s^2 3p^6$

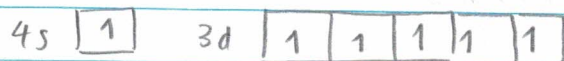
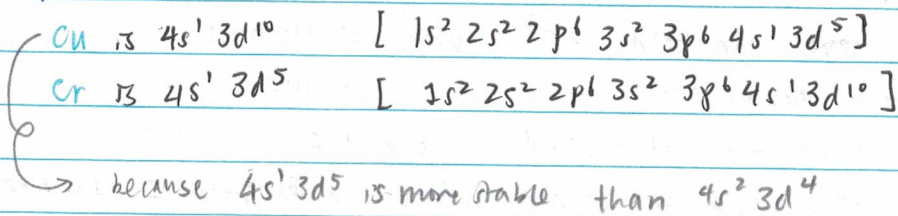
shorthand = nearest noble gas configuration.

e.g. Zr is $[Kr] 5s^2 4d^2$

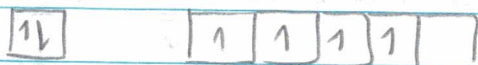
Cs is $[Xe] 6s^1$



exceptions



vs



Electronic configuration of IONS

- electrons furthest from the nucleus are lost first
- 4s are lost before 3d (4s gains energy when occupied)
- electrons are removed in numerical order (4 before 3, 3 before 2, etc)
- isoelectronic: same electronic configuration.

12.1 Electrons IN ATOMS

evidence for the existence of sub-shells & shells can be seen from studies in trends of ionisation energies within the periodic table.

first ionisation energy: energy absorbed when 1 MOLE of electrons is removed from 1 MOLE of an atoms in GASEOUS state, giving 1 MOLE of positive ions.



ionisation energy (1st) is different, depending on...

① **DISTANCE**: → greater the distance between nucleus + valence electrons
→ less attractive force
→ decrease in IE

② **SHIELDING**: → repulsion filled by inner shells/subshells makes the electrons in the outer shell easier to remove
→ decrease in IE.

③ **NUCLEAR CHARGE**: → nucleus is +ve charge
→ more protons, more charge, more pull (force of attraction)
→ increase in IE

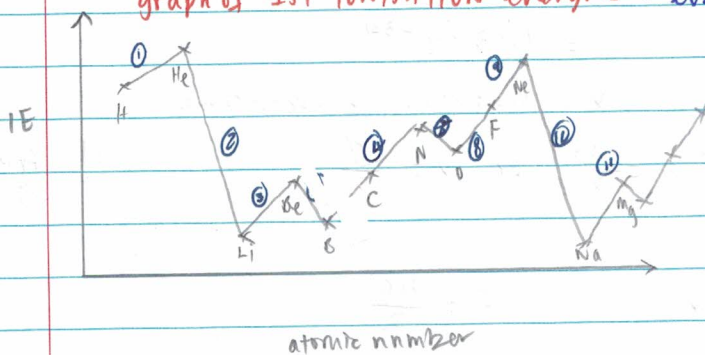
④ **ELECTRON PAIRING**: → paired e^- is easier to remove than unpaired e^-
→ electrons in shells that are half full are hardest to remove
e.g. $3p^3$

1	1	1
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successive ionisation energies: evidence for SHELLS.

↳ after removing FIRST electron, can remove other electrons.

graph of 1ST IONISATION energies: evidence for SUBSHELLS.



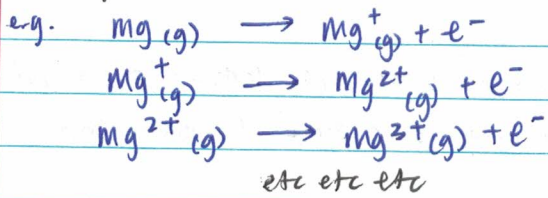
1. $H \rightarrow He$: bigger nucleus (+ve charge), increase
2. $He \rightarrow Li$: distance from nucleus (shell) ↓ IE
· increase in shielding $Be \rightarrow B$: new shell ↓ IE
3. $Li \rightarrow Be$: increased nuclear charge ↑ IE
4. $B \rightarrow C \rightarrow N$: increased nuclear charge ↑ IE
5. $N \rightarrow O$: in $N(2p^3)$, all unpaired, in $2p^4$ increased repulsion between paired electrons = ↓ IE
6. $O \rightarrow F \rightarrow Ne$: increased nuclear charge ↑ IE

EVIDENCE for SHELLS:

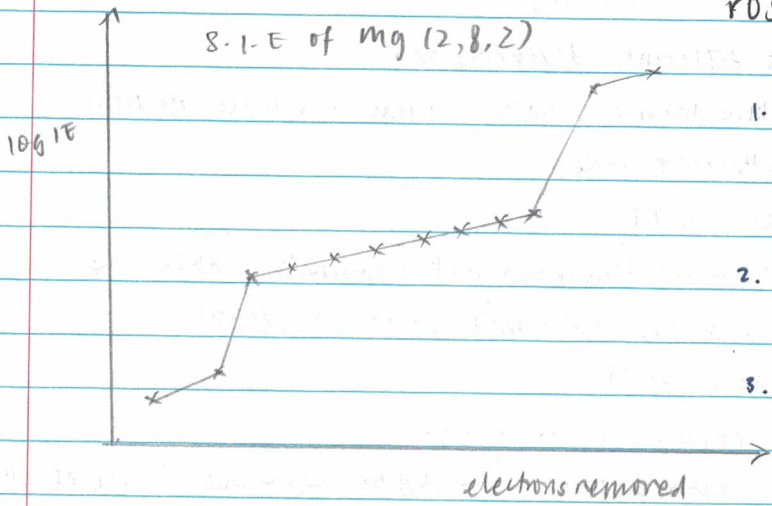
→ large decrease in IE from group 18 to group 1 charge (since all e^- removed same shell)
due to an increased distance of the outer electron from the nucleus when a new shell is added ($He \rightarrow Li$, $Ne \rightarrow Na$)

SUCCESSIVE IONISATION ENERGIES. (evidence for shells!)

you can continue to remove electrons (2nd, 3rd... ionisation energies.)



} There is a general increase
 ① because ratio of protons: electrons is bigger = greater attraction
 ② removing electrons forms a GREATER POSITIVE CHARGE.



1. large increase - 2nd electron removed from 3rd shell, 3rd one is removed from 2nd \rightarrow closer to nucleus.
2. general increase = higher ratio of +ve: -ve
3. large increase - 10th electron from 2nd shell, 11th from 1st = even closer to nucleus (closer than 2nd)

• the FIRST large INCREASE indicates no. of electrons in the outer shell.

QUANTIZATION OF ENERGY

precise lines in line emission spectrum have a specific wavelength which corresponds to a discrete amt of energy (= energy difference between energy levels)

$\rightarrow \therefore$ it is foundation of quantization: radio electromagnetic radiation comes in parcels / discrete.

\rightarrow photon is a quantum of radiation. [frequency, energy, related by...]

$E = h\nu$ $h = 6.63 \times 10^{-34}$
 $\nu =$ frequency of radiation.

[frequency + wavelength related by...]
 $c = \nu\lambda$ or $\nu = \frac{c}{\lambda}$!! frequency is $\frac{c}{\lambda}$

therefore $E = h\nu$ is $E = \frac{hc}{\lambda}$

$E \propto \frac{1}{\lambda}$ (greater energy, smaller wavelength, higher frequency)

EXAM QUESTIONS & EXAMPLES

examples: photon of red light, 4.5×10^{14} s⁻¹ — $\lambda = 650 \text{ nm}$, $h = 6.626 \times 10^{-34}$, $c = 2.998 \times 10^8$

find the energy.

$$E = \frac{hc}{\lambda} \Rightarrow \frac{6.626 \times 10^{-34} \times 2.998 \times 10^8}{650 \times 10^{-9}} = 3.056 \times 10^{-19}$$

* wavelength must always be in m.

first ionisation energy in kJ mol^{-1} — $\lambda = 91.16 \text{ nm}$, $h = 6.626 \times 10^{-34}$, $c = 2.998 \times 10^8$
 $6.626 \times 10^{-34} \times 2.998 \times 10^8$ $N_A = 6.022 \times 10^{23}$

$$\begin{aligned} E &= \frac{hc}{\lambda} = \frac{6.626 \times 10^{-34} \times 2.998 \times 10^8}{91.16 \times 10^{-9}} \\ &= 2.179 \times 10^{-18} \text{ per photon} \\ &= 2.179 \times 10^{-18} \times 6.022 \times 10^{23} \text{ per mole} \\ &= 2.179 \times 10^{-18} \times 6.022 \times 10^{23} \times 10^3 \text{ per kJ/mol} \end{aligned}$$

first ionisation energy in kJ mol^{-1} , frequency is 3.3 PHz ,

$$\star c = f\lambda \text{ so } f = \frac{c}{\lambda} \text{ and } \frac{c}{f} = \lambda$$

$$\frac{2.998 \times 10^8}{3.3 \times 10^{15}} = \lambda \quad \dots \text{ and work out the rest.}$$

state a radioactive isotope used in medicine.

• iodine - 131.

bromine exists as the isotopes Br-79 and Br-81. What is % of Br-79 with 79.9 ma

$$79x + 81(100-x) = 79.90$$

$$-2x = -110, \quad x = 55\%$$

the value of the relative molecular mass of HBr is 80.91. why doesn't any sample of HBr have this Mr value? (2)

① it is an average of all HBr values

② each HBr has a different mass depending on the isotopes of H and Br.

describe and explain how the chemical/physical properties of 2 isotopes compare.

① similar chemical properties. ② same valence electrons

③ different physical properties ④ different masses.

explain how the formation of lines in a visible line spectrum indicates the presence of energy levels. (1)

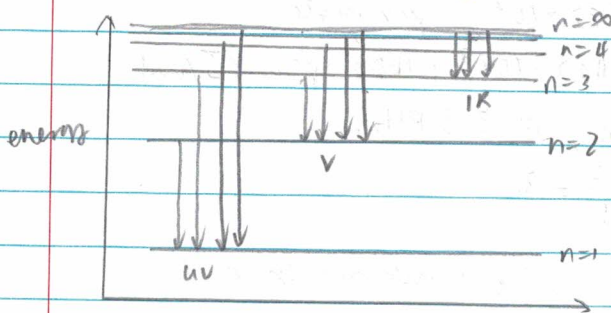
• shows electrons transition from high to low energy levels, and so they emit energy in the form of light

describe the emission spectrum of hydrogen; explain how this is related to energy levels.

- ① emission spectrum is a series of lines that converge @ higher frequencies
 - ② electrons transfer from high to low energy levels and light is emitted.
 - ③ as they go down the energy levels (releasing energy)
 - ④ $n=1$ is ground = UV, $n=2$, visible light, $n=3$, IR. transitions to $n=1, 2, 3$.
- Ryman
Balmer
Paschen

the line which represents the FIRST IONISATION ENERGY of hydrogen is from $n=1$ to $n=\infty$ (to remove / ionise, electron must be taken OUT)

energy level diagram for hydrogen



what is meant by the term "first ionisation energy"?

the amount of energy absorbed when one more of electrons is removed from one more of ~~the~~ a gaseous atom.

what is "periodicity"?

repeating patterns / trends across periods.

explain how the graph of first ionisation energies provides evidence for the existence of main energy levels and sub-levels

- existence of shells - huge drop between He and Li (different shells)
- existence of subshells - general increase across a period. (electrons added to same shell)
 - ↳ furthermore = highest value for noble gases, lowest for alkali
 - ↳ drop in IE from Be to B (s^2 to s^2p^1) and drop in IE from N to O (p^3 to p^4)
 - ↳ value drops drastically after Gr 8 as a new shell added = further away from nucleus.
 - ↳ greater attraction from higher z_{eff} ratio.

Why is the difference between Germanium's 4th and 5th ionization energies

is much greater than other successive values?

instead of $n=4$ (taking the electron) taken from $n=2$ (taken from energy level closer to the nucleus)

↳ attraction of protons is greater

EXAM QUESTIONS

define the term isotope.

atoms of the same element but with different NUMBER OF NEUTRONS (not mass)
state how a line spectra differs from a continuous spectrum.

line spectrum = discrete LINES at certain frequencies.

continuous spectrum = all possible wavelengths/frequencies

successive ionization energies:

1st - 799. 2nd - 2420 3rd - 3660 4th - 25000 5th - 32800

↳ why is there a large increase between 3 → 4?

electron taken from $n=1$ level instead of $n=2$ = closer to nucleus

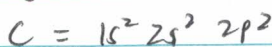
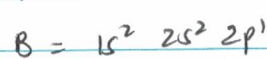
more attraction = more energy needed to ionise.

↳ why is the increase between 1st → 2nd > 2nd → 3rd?

electron in 1 → 2 taken away from a different shell; closer to

nucleus (2p to 2s) ⇒ more attraction ↑ IE.

↳ state with reason how the value for the second IE of carbon would compare with the second IE of boron.



} carbon = less IE because taking away from $2p^2$,
further away than $2s^2$ → less attraction = less IE.
whereas B taken from 2s level = closer = ↑ IE.

* provide 2 sides!!

However, nuclear charge of C is greater. ↑ IE due to ↑ protons than Boron (tve: -ve ratio higher).

technitium is used as a tracer in medicine. suggest why it could be dangerous

radioactive isotope which could ionise body's cells = cause cancer.

what is relative atomic mass?

the weighted mean mass of all atoms and isotopes (naturally occurring)

relative to a C-12 atom ($1/12$ th)

why is there a general increase in the value of 1st IE Li → Ne

all in the same shell, relatively the same distance from the nucleus ($n=2$ energy level), hence a general increase merely based on nuclear attraction (tve: -ve ratio) as proton no. increase as atomic no. increases.

[ATOMIC RADIUS DECREASES]

first ionization energy of hydrogen is 1312 kJ mol^{-1} . determine frequency and wavelength.

$$E = h\nu \leftarrow \text{frequency}$$

↑ planck's constant

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

explain how the line spectrum is related to the energy level in a hydrogen atom.

- ↳ electrons jump from excited state to the $n=2$ level
- ↳ converge @ higher frequencies
- ↳ frequencies of the light = directly proportional to the difference in energy levels.