Lesson 7 For Book 1
Review Exerice Drawing of unit cell
The lattice structure of BaO (s) is described as the interpenetration of two simple
cubic lattices, one of $Ba^{2+}$ ions and the other of $O^{2-}$ ions.
a) Draw the unit cell of BaO (s), labelling the $Ba^{2+}$ and $O^{2-}$ ions.
(Hint = use a ruler to draw the cubic first.)
b) What is the coordination number of each $O^{2-}$ ion in the structure?
$\rightarrow$ You should remember the number (Lesson 6) unless you can find it by looking at
what you have drawn. (for the anion only!) (8)
Definition of Radii
1. <b>Ionic radii</b> = the radii of an perfect ionic compound, that is, the internuclear
distance $\rightarrow$ the sum of the cationic and anionic radii.
2. <b>Covalent radii</b> = half the internuclear distance between two atoms in a c
bonded molecule
$\rightarrow$ half of the bond length of <b>homoatomic</b> covalent molecules e.g.
$H_2, O_2$
$\rightarrow$ the sum of the covalent radii of atom A and B (heteroatomic)
3. Metallic radii = half the internuclear distance between two metallic atoms/ ions
in a metallic crystal.
4. <b>van der Waal's radii</b> = half the distance between the nuclei of two atoms in
adjcent molecules
$\rightarrow$ You should bear in mind that the consideration of "radii" can help us compare
some physical properties of different compounds e.g. strength of all bonds, m.p./b.p.
of all compounds, electrical conductivity of m, polarizability of a,
polarizing power of c
Exercise 1 An iodine molecule can be represented but he diagram below, with each
dot representing an atomic nucleus.
i) Using one or more diagrams of this kind, illustrate your
understanding of the two terms "covalent radius" Rc and
"van der Waal's radius" Rv.
ii) Which of the values (Rc or Rv) is larger than the another? Account for your
anwser.
$\rightarrow$ Be careful, you should be clear of the concept of covalent bond (wa molecule)

and intermolecular force (**b**\_\_\_\_\_\_ two molecules).

Different kinds of compounds
1. Molecular structure = S molecular and M structure
$\rightarrow$ depends on molecule's size e.g. <b>Iodine</b> is s molecule while some <b>plastic</b> is
m molecule as plastic consists of a large c
а с-я-с + анды-я-цы, - с в с к я д · + 2 ндо но он - к - с в с к я д · + 2 ндо
iodine in Face centred cubic General condensation polymerization
reaction for nylon
$\rightarrow$ They are made up of <b>discrete</b> molecules held by inter force
(van force and h bond).
$\rightarrow$ As for simple molecules, they have b.p. / m.p. as the intermolecular force is
w They can conduct electricity as they have <b>no</b> delocalized electrons /
m ions.
$\rightarrow$ The existence of H bond will increase the boiling / melting point.
Exercise 2 Comparison of boiling point
a) Draw the Lewis structure of a $H_2O$ molecule and $F_2O$ molecule.
(Hint = Count the no. of valence electrons)
b) Which compound, $H_2O$ or $F_2O$ , would have a higher b.p.? Explain your anwser.
2. Giant Structure
$\rightarrow$ all the atoms or ions present in the lattice are linked by strong bonding.
$\rightarrow$ Giant Ionic (Lesson 6), Giant covalent and Giant metallic (Lesson 6)
Giant Covalent Compounds
Giant Covalent structure is one in which all atoms are linked together by a
continuous system of well-defined electron-pair bonds (c / /
bonds). The bonds are thus <b>diectional</b> .
$\rightarrow$ High melting and boiling point
Example Allotropes of carbon
a is d; b is g;
d is $C_{60}$ and e is $C_{540}$ ;
$\rightarrow$ Which one/two of the allotropes has/have the Lowest m.p?
and as they are just <b>discrete molecules</b> .!

Exercise 3 Something about hydrides

Consider the hydrides of three period 3 elements: SiH<sub>4</sub>, PH<sub>3</sub> and H<sub>2</sub>S.

For PH<sub>3</sub>

a) For each hydride, draw a 3-D structure showing the bond electron pairs and lone electron pair (s), if any, of the central atom.

<u>For  $H_2S$ </u>

Valence electron = 8

central lone pair electron = 0

b) Explain why  $H_2S$  has a higher boiling point.

(**hint** = Actually, they are all simple molecule. The difference in b.p. is not due to their types of structure.)

Extra Concept

1) Solubility of compounds in some solvent

 $\rightarrow$  In Lesson 5, you should learn that what solubility depends on, i.e. intermolecular force.

 $\rightarrow$  In detail, Solubility is the property of a s\_\_\_\_, l\_\_\_ or g\_\_\_\_ chemical

substance (i.e. **solute**) to dissolve in a solid, liquid, or gaseous **solvent** to form a homogeneous solution of the solute in the solvent

 $\rightarrow$  How well the solute can mix with the s\_\_\_\_\_

→ It depends on the use of solvent (p\_\_\_\_\_ or non p\_\_\_\_) and the kinds of solute (polar or non polar)

## <u>Like-disolve-like concept</u>

To dissolve a solute, we need to break the s\_\_\_\_\_ and s\_\_\_\_\_ interactions (I<sub>1</sub>). Then, we have the new formation of s\_\_\_\_\_ and s\_\_\_\_\_ interactions (I<sub>2</sub>).

 $\rightarrow$  If I<sub>2</sub> is stronger / more exo\_\_\_\_\_ than I<sub>1</sub>, the solute is said to be soluble in that solvent.

 $\rightarrow$  In fact, polar solute likes polar solvent while \_\_\_\_\_ polar solute like non polar solvent as the type of interactions involved should be similar.

→ We should know that **organic solute** can "only" be dissolved by \_\_\_\_\_\_ solvent. <u>New concept --- For ionic compound</u>

As for ionic compounds and water as solvent, to dissolve them, we need to break the i\_\_\_\_\_ lattice. What is forming is the electrostatic attraction between p\_\_\_\_ water molecule and those m\_\_\_\_\_ ions. Thus, we have,

 $\Delta H_{\text{soln}} = \Delta H_{\text{hydration}} - \Delta H_{\text{lattice}} \rightarrow \text{Energetical aspect!}$ 

Exercise 4 Determination of solubility

Try to arrange the solubility of  $CO_2$ ,  $NH_3$  and  $O_2$  in water in descending order with explanation.  $(NH_3 > CO_2 > O_2)$ 

2) The value and the sign of  $\Delta H_{\text{lattice}}$  of ionic compounds

 $\Delta H_{\text{lattice}}$  measures the energetical stability of an I\_\_\_\_ crystal/salt, which is formed by the combination of an a\_\_\_\_\_ and a cation.

 $\rightarrow$  if the sign is negative, the ionic crystal is energetically stable.

## Soft ions and Hard ions

• Soft ions = those ions with high r\_\_\_\_ to charge ratio, i.e., have a large electron cloud and have a high polarizability

e.g. Rb+, I-

• **Hard ions** = those ions with low radius to c\_\_\_\_\_ ratio, i.e. have a small ionic Size and have a high polarizing power.

e.g. Li+ , F-

**Concept** = Soft ions like s\_\_\_\_\_ ions while hard ions like h\_\_\_\_\_ ions

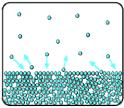
 $\rightarrow$  Similar ions with similar size will combine to form a more stable ionic compound as they can have a better overlapping/ packing in lattice.

**soft cations +soft anions** / **hard cations +hard anions** can form a more stable ionic salt!

## Vapour pressure of a solution

All liquid will evaporate to form vapour at all temperature (except at absolute zero) as the molecules are gaining K.E. . The term vapour pressure relates to the tendency of particles to escape from the liquid.

→ If a liquid has a high tendency to evaporate, the liquid is said to be V\_\_\_\_.



 $\rightarrow$  Hence, the liquid has a high/low ? vapour pressure. (at a certain temp.)

Look at the diagram, you should realise that if a liquid is volatile, there are more molecules exerting a force to the container  $\rightarrow$  higher vapour pressure.

**Concept** = A liquid with high vapor pressure will have a lower/higher boiling point. *Extra Knowledge --- Ionic Fluid* 

- An ionic liquid (IL) is a **salt** in the liquid state. They can be used as powerful **solvents** and **electrically conducting** fluids (i.e. electrolytes).
- Ionic liquids are often moderate to poor conductors of electricity

   (as they have \_\_\_\_ mobile ions), non-ionizing (e.g. non-polar), highly
   viscous and frequently exhibit low vapor pressure (Why? As the bond involved is \_\_\_\_ bond!).