CHEMICAL BONDING TUTORIAL

Interatomic Bonding

1 Using the list below, give one example to each of the following solid lattices showing the given properties. Name the structure and type of bonding found in each lattice.

graphite, diamond, copper, iodine, sodium chloride

- (a) A hard and brittle solid.
- (b) A solid that conducts electricity and melts at 1085°C.
- (c) A solid that cuts through metal and does not conduct electricity.
- (d) A substance that sublimates when heated.
- (e) A solid that can be used as a lubricant.

	Structure	Bonding type	Example
(a)	ionic lattice	Ionic bonding	NaC <i>l</i>
(b)	metallic lattice	Metallic bonding	Cu
(c)	Giant molecular structure	Covalent bonding	Diamond
(d)	Simple molecular structure	 Covalent bonding within molecule Weak intermolecular force of attraction between molecules 	I ₂
(e)	Giant molecular structure	 Covalent bonding between atoms within a layer Weak intermolecular force of attraction between layers 	Graphite

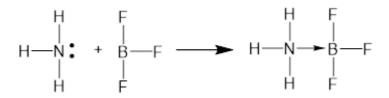
2 Which statement(s) are correct?

- I HCl (g) can conduct electricity.
- II Covalent bond is a weak bond compared to ionic bond as boiling point of H₂O is lower than that of MgO.
- III Ionic bonds and covalent bonds can both occur in the same compound.
- **IV** Metals can be distinguished from ionic compounds by their electrical conductivity in the solid and liquid state.
- A Statements I and III are correct.
- **B** Statements **II** and **IV** are correct.
- C Statement III and IV are correct.
- D Statements I, III and IV are correct
- False. HC/(g) has simple molecular structure with no mobile charge carrier. HC/can dissociate into ions H⁺(aq) and Cl⁻(aq) in water, and conduct electricity in aq state.
- II False. Both are strong bonds. The boiling of H₂O involves only the breaking of the weak hydrogen bonds between the H₂O molecules. However, the strong ionic bond is overcome in MgO when it undergoes phase change during boiling point.

- III True. Eg. Na₂SO₄. Ionic bonds between Na⁺ and SO₄²⁻ ions, covalent bonds between S and O in SO₄²⁻ ion.
- **IV** True. Metals conduct electricity in molten and solid state, while ionic compounds conduct electricity in molten state, not in solid state. Hence, this can be used to distinguish metal from ionic compounds.
- **3** Boron trifluoride and ammonia reacts in a 1:1 mole ratio to form a single compound. Describe the type of bond formed during this reaction.

N atom in ammonia forms a dative bond to B atom in BF₃.

B atom in BF_3 is electron deficient, hence B attains the octet by accepting a lone pair from N in ammonia into its empty p-orbital.



- 4 Explain the following observations in terms of structure and bonding.
 - i. MgO has a higher melting point than NaCl.

Both MgO and NaCl have ionic lattice structures.

- Product of charges (q₊ × q₋) is larger for MgO and interionic distance (r₊ + r₋) is smaller for MgO.
- Since $|L.E| \propto \left| \frac{q_+ \times q_-}{r_+ + r_-} \right|$, magnitude of lattice energy of MgO is greater than NaCl.
- More energy is needed to overcome the stronger ionic bonds in MgO than in NaCl.
- MgO has a higher mp than NaCl.
- ii. At room temperature and pressure, CO₂ is a gas, while SiO₂ is a solid of high melting temperature.

 CO_2 exists as simple molecular structure with weak intermolecular forces/ instantaneous dipole - induced dipole (id-id) interactions between CO_2 molecules. SiO₂ exists as a giant molecular structure whereby all atoms are bonded to each other by strong covalent bonds.

The energy supplied at room temperature is sufficient to overcome weak intermolecular forces between CO_2 molecules to allow CO_2 to exist as a gas.

However, energy is insufficient to overcome the strong covalent bonds in SiO_2 hence, SiO_2 exist as solids at room temperature.

Note: must relate back to physical states of the compounds at rtp. The question is not focusing about the mp/bp.

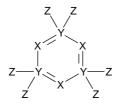
Dot-and-Cross Diagram and VSEPR

5 Fill up the table below for the following compounds:

	compound	'dot-and-cross' diagram	 (i) Draw and name the shape of the covalent molecule or polyatomic ion (ii) Indicate the bond angle in the diagram 	Bonds or intermolecular forces broken during melting or boiling process
(i)	Al ₂ O ₃ (m.p. 2072°C) Ionic compound	$2\left[Al\right]^{3+}3\left[\overset{\times}{\cdot}\overset{\times}{\overset{\times}{\Omega}}\overset{\times}{\overset{\times}{\Omega}}\right]^{2-}$	NA	Ionic bond
(ii)	A <i>l</i> C <i>l</i> ₃ (m.p. 192°C) Covalent molecule	:C <i>l</i> : :C <i>l</i> -×Å <i>l</i> וC <i>l</i> :	CI CI CI Trigonal planar 120°	Instantaneous dipole-induced dipole
(iii)	PC <i>l</i> ₃ (I)	:C/•× P וC/: :C/:	Cl 107° Cl trigonal pyramidal, 107°	Permanent dipole-permanent dipole & Id-id
(iv)	CH ₂ C <i>l</i> ₂ (I)	H : Ċl··× Č H	H 109.5° H C''''Cl Cl tetrahedral, 109.5°	Permanent dipole-permanent dipole & Id-id
(v)	PCl₅(s)	:C/: :C/·*P.*C/: :C/:C/:	CI 90° CI CI CI CI CI CI CI CI CI CI	Instantaneous dipole-induced dipole
(vi)	HCN	HוC:×N ×	$H \xrightarrow{C} C \xrightarrow{C} N$ Linear	Permanent dipole-permanent dipole & Id-id

compound	'dot-and-cross' diagram	 (i) Draw and name the shape of the covalent molecule or polyatomic ion (ii) Indicate the bond angle in the diagram 	Bonds or intermolecular forces broken during melting or boiling process
(vii) SO ₂	Ö:* Š *:Ö	Bent, 118º (<120º)	Permanent dipole-permanent dipole & Id-id
(viii) SO3	:0: Ö:* Š ::Ö	O O Trigonal planar 120°	Instantaneous dipole-induced dipole
(ix) H ₂ O ₂ (H-O-O-H arrangement)	H•× over the term of term	H around each O Bent 105°	Hydrogen Bond & Id–id
(x) (NH4)2SO4	$2 \begin{bmatrix} H \\ H \cdot \cdot \cdot N \\ H \cdot \cdot N \\ H \end{bmatrix}^{+}$ $\begin{bmatrix} \cdot & \cdot \\ H \\ H \end{bmatrix}^{2-}$ $\begin{bmatrix} \cdot & \cdot \\ \vdots \\$	$2 \begin{bmatrix} H \\ H \\ H \\ H \end{bmatrix}^{+}$ tetrahedral about N and S $\left[\begin{array}{c} 0 \\ H \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\$	lonic bond

6 A stable molecule containing atoms of the elements, X, Y and Z has the following structure.

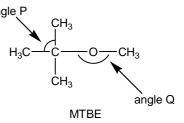


Which option is a possible combination of the elements?

	<u>X</u>	<u>Y</u>	<u>Z</u>
A	N	P	C/
В	0	S	C <i>l</i>
С	В	N	Н
D	Р	0	F

- Y formed 5 covalent bonds → Group 15, Period 3 and beyond
- X formed 3 covalent bonds. Could be Group 13 or 15.
- Z formed 1 covalent bond → Either hydrogen or Group 17
- 7 MTBE is a constituent of petrol. What are the values of angle P and angle Q in a molecule of MTBE? $H_3C - CH_3$

	angle P	angle Q
Α	90°	105°
В	90°	180°
С	109°	105°
D	109°	180°



Answer C

- Central atom, C, has 4 bond pairs and no lone pair → Tetrahedral in shape with bond angle of 109.5°
- Central atom, **O**, has 2 bond pairs and 2 lone pair → Bent in shape with bond angle of 105°
- **8** Using VSEPR theory, predict which compound in each of the following pair of molecules has a larger bond angle.
 - (a) BCl_3 and NCl_3 (b) H_2S and PH_3
 - (a) BC^{*l*}_b has 3 electron regions (3 bond pairs) whereas NC^{*l*}_b has 4 electron regions (3 bond pairs and 1 lone pair).

By VSEPR theory, the electron regions around the central atom arrange themselves to be as far apart as possible to minimise mutual repulsion.

Therefore, BC l_3 has a trigonal planar shape and the bond angle is 120 ° whereas NC l_3 has a trigonal pyramidal shape and the bond angle is 107 °.

BCI₃ has a larger bond angle.

(b) H₂S and PH₃ has 4 electron regions each. H₂S has 2 bond pairs and 2 lone pairs whereas PH₃ has 3 bond pairs and 1 lone pair.
 By VSEPR theory, lone pair-lone pair repulsion is stronger than lone pair-bond pair repulsion which is in turn stronger than bond pair-bond pair repulsion.
 In H₂S, the stronger lone pair-lone pair repulsion forces the bond pairs to be closer together,

In H₂S, the stronger ione pair-ione pair repulsion forces the bond pairs to be closer together, decreasing the bond angle to 105° as compared to 107° in PH₃.

Therefore, PH_3 has a larger bond angle.

9 [N10/III/4c]

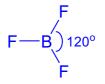
(a) Draw 'dot-and-cross' diagrams to show the bonding in the molecules of NO₂, O₃ and BF₃.

In the molecule NO₂, the central atom is nitrogen. In each case you should distinguish carefully between electrons originating from the central atom and those from the two outermost atoms. Include all lone pairs in your diagrams.



(b) Suggest a value for the bond angles in each of the three molecules, giving reasons for your choice.

 BF_3 – central atom has 3 bond pairs and 0 lone pair. Therefore, BF_3 has a trigonal planar shape and the bond angle is 120 °.



 O_3 – central atom has 2 bond pairs and 1 lone pair. Since lone pair-bond pair repulsion > bond pair-bond pair repulsion , estimated angle = 118 ° (slightly less than 120°).



 NO_2 – central N atom has 2 bond pairs (1 single covalent bond and 1 double bond) and 1 unpaired electron. The unpaired electron cause less repulsion than a lone pair so estimated angle = 130 ° (Note: accept angles greater than angle given for O₃ but less than 180 °, assuming that the repulsion of unpaired electron is less than the repulsion of bond pair (2e⁻))



Polarity and Intermolecular Forces

10 (a) Classify the following molecules as polar or non-polar.

 $NH_3,\,SF_6,\,CH_2 = CH_2,\,CH_3OH,\,C/F_3,\,CH_3OCH_3\,,\,N_2H_4$

Non-polar	Polar
SF ₆ , CH ₂ =CH ₂	NH ₃ , CH ₃ OH, CH ₃ OCH ₃ , C/F ₃ N ₂ H ₄

(b) Hence, classify the molecules in (a) according to the main type of intermolecular forces present.

Instantaneous dipole- induced dipole	Permanent dipole- permanent dipole	Hydrogen bonds
SF ₆ , CH ₂ =CH ₂	CH_3OCH_3, ClF_3	NH ₃ , CH ₃ OH, N ₂ H ₄

Structure and Physical Properties

11 The boiling points of four compounds are given in the table below.

Compound	Boiling point/ °C
H ₂ O	100
CH₃OH	65
SiH ₄	-107
CH ₄	-164

Explain the differences in boiling point between

(a) CH₄ and SiH₄

Both CH₄ and SiH₄ are covalent compounds with simple molecular structures. They are nonpolar and have instantaneous dipole-induced dipole (id-id) interactions between molecules. SiH₄ has a larger electron cloud than CH₄ which is more easily distorted resulting in stronger id-id interactions.

More energy is required to overcome the stronger id-id interactions between SiH_4 molecules than those between CH_4 molecules.

Therefore, SiH₄ has a higher boiling point than CH₄.

(b) H₂O and CH₃OH

Both H_2O and CH_3OH are covalent compounds with simple molecular structures. They are polar with H-bonds between molecules.

 H_2O have more extensive H-bonds (average 2 per molecule) than CH_3OH (average 1 per molecule).

More energy is required to overcome the more extensive H-bonds between H_2O molecules. Therefore, H_2O has a higher boiling point than CH_3OH . **12** The boiling points of three organic compounds are given in the table below.

Compound	Molecular formula	<i>M</i> r	Boiling point/°C
Α	CH ₃ CH ₂ OCH ₂ CH ₃	74	35
В	CH ₃ CH ₂ CH ₂ CH ₂ OH	74	118
С	(CH₃)₃COH	74	83

Explain the differences in boiling point between

(a) A and B

Both are covalent compounds with simple molecular structures. They are polar molecules with the same $M_{\rm r}$.

Less energy is required to overcome the weaker permanent dipole-permanent dipole interactions between $CH_3CH_2OCH_2CH_3$ than the stronger H-bonds between $CH_3CH_2CH_2CH_2CH_2OH$ molecules

Therefore, compound A has a lower boiling point than compound B.

(b) B and C

Both are covalent compounds with simple molecular structures. They are polar molecules with the same M_r . Both compounds have H-bonding and instantaneous dipole-induced dipole (id-id) interaction between the molecules.

CH₃CH₂CH₂CH₂OH is a straight chain molecule and it has a larger surface area of contact, as compared to branched chain (CH₃)₃COH. More energy is needed to overcome the stronger id-id interactions between CH₃CH₂CH₂CH₂OH molecules than those between (CH₃)₃COH molecules. Therefore, CH₃CH₂CH₂CH₂OH has a higher boiling point than (CH₃)₃COH.

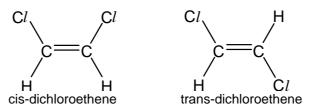
(Note: strengths of H-bond in $CH_3CH_2CH_2CH_2OH$ and in $(CH_3)_3COH$ are similar, so students have to recognise that it is the id-id (due to different surface area of contact) that explains their difference in boiling points.)

- **13** Referring to table in question **5**, explain the following observations.
 - (i) Al₂O₃ conducts electricity in molten state while AlCl₃ in molten state does not.
 Al₂O₃ has an ionic lattice structure, while and AlCl₃ has a simple molecular structure.
 In molten state, Al³⁺ and O²⁻ ions in Al₂O₃ act as mobile charge carrier to conduct electricity.
 There is no mobile charge carrier in AlCl₃ molecules, hence it cannot conduct electricity in molten state.
 - (ii) Difference in physical state of PCl_3 and PCl_5 .

Both PCl_3 and PCl_5 have simple molecular structure.

- There is permanent dipole-permanent dipole (pd-pd) and instantaneous dipole-induced dipole (id-id) interaction between PCl₃ molecules, while there is id-id between PCl₅ molecules.
- PCl₅ has a larger electron cloud that can be more easily distorted, to give a stronger id-id interaction between PCl₅ molecules.
- At room temperature, there is sufficient energy to overcome pd-pd & id-id interactions between PCl₃ molecules, but not the stronger id-id interaction between PCl₅ molecules.
- Therefore, PCl₅ exists as solid while PCl₃ exists as liquid.

14 (a) The boiling point of cis-dichloroethene is 333 K, whereas that of trans-dichloroethene is 321 K.



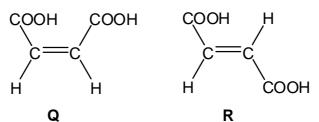
In cis-dichloroethene, the bond dipole moments are not cancelled out. Cis-dichloroethene has a net dipole moment and it is polar.

In trans-dichloroethene, the bond dipole moments are cancelled out. Trans-dichloroethene has zero net dipole moment and it is non-polar.

More energy is needed to overcome the stronger Permanent dipole-permanent dipole and instantaneous dipole-induced dipole (id-id) interactions between cis-dichloroethene molecules than the id-id interactions between trans-dichloroethene molecules.

Therefore, cis-dichloroethene has a higher boiling point than trans-dichloroethene.

(b) The boiling point of **R** is higher than **Q**.



In **Q**, due to the close proximity of the 2 COOH groups, intramolecular H-bonds are formed, leaving fewer sites for intermolecular H-bonding. R forms only intermolecular H-bonds.

More energy is required to overcome the **more extensive intermolecular H-bonding between R molecules** than between **Q** molecules. Therefore, **R** has a higher boiling point than **Q**.

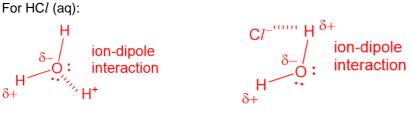
Solubility

- 15 Predict with explanations whether or not
 - (a) CH₃CH₂OH is soluble in H₂O Energy given out in the formation of H-bonds between CH₃CH₂OH and H₂O molecules is sufficient to compensate for the energy taken in to overcome the H-bonds between CH₃CH₂OH molecules and those between H₂O molecules. Therefore, CH₃CH₂OH is soluble in H₂O.
 - (b) NH₄NO₃ is soluble in benzene, C₆H₆. NH₄NO₃ does not dissociate to give ions in a non-polar solvent (C₆H₆). No favourable interaction is possible between NH₄NO₃ and C₆H₆. There is insufficient energy to overcome the strong ionic bonds between NH₄⁺ and NO₃⁻ and instantaneous dipole-induced dipole interactions between C₆H₆. Therefore, NH₄NO₃ is insoluble in C₆H₆.

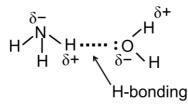
16 Ammonia and hydrogen chloride gases are soluble in water because they interact with the solvent.

 $\begin{array}{rcl} \mathsf{HC}l(\mathsf{g}) + \mathsf{aq} & \to & \mathsf{H^+}\left(\mathsf{aq}\right) + \mathsf{C}l^-\left(\mathsf{aq}\right) \\ \mathsf{NH}_3(\mathsf{g}) + \mathsf{aq} & \to & \mathsf{NH}_3\left(\mathsf{aq}\right) \end{array}$

Use suitable diagrams to illustrate all the possible interactions between the dissolved gas and **a water** molecule.



For NH₃(aq) :



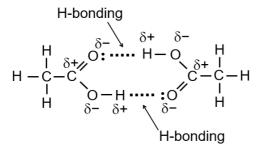
Application questions

- 17 Provide an explanation for the following observations.
 - (a) Ice is less dense than water.

Due to the formation of four permanent H-bonds per H₂O molecule in ice, ice has a more open structure than liquid water. Ice occupies a larger volume than water for the same number of molecules. Therefore, ice is less dense than water.

(b) The relative molecular mass of ethanoic acid is 120 in benzene, C₆H₆.

In a non-polar solvent like benzene, ethanoic acid dimerises by forming H-bonds between 2 molecules. As a result, the apparent M_r is double of 60.

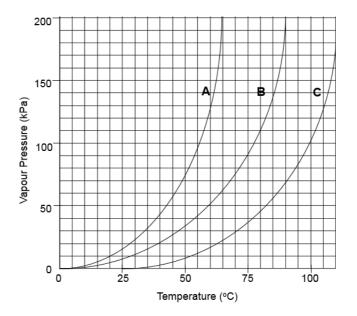


18 Vaporization occurs when some molecules in a liquid possesses enough kinetic energy to escape from the surface of the liquid at a given temperature.

Saturated vapour pressure is the pressure on the walls of the container exerted by the gas molecules vaporized from the surface of the liquid in a closed container when the rates of condensation and evaporation are equal.

Boiling occurs when the vapour pressure is equal to the external pressure which is usually the atmospheric pressure (101 kPa).

The graph below shows the vapour pressures of three liquids at varying temperatures.



(a) The identities for liquids A and B could be ethanol (CH₃CH₂OH) and propanone (CH₃COCH₃). Based on the graph, identify liquids A and B and explain your reasoning. [4]
 A: propanone [1]. B: ethanol [1]

At any given temperature, **A** has a higher vapour pressure than B (**A** has a lower boiling point than B) [½], showing that the intermolecular forces in A are weaker than that in B.

Permanent dipole-permanent dipole interactions between propanone molecules [½] is weaker [½] than the hydrogen bonds between ethanol molecules [½]. [correct identification and comparison of strength of IMF] At any given temperature, more propanone molecules have enough kinetic energy to overcome the intermolecular forces and exist as vapour, resulting in a higher vapour pressure. Therefore, A is propanone and B is ethanol.

(b) Suggest a possible identity for C. H₂O, boiling point \approx 100 °C (Temp = 100 °C when pressure = 101 kPa)