# Q1.

Which statement about intermolecular forces is not correct?

	Α	Intermolecular forces exist between all simple molecules.	0	
	В	Hydrogen bonding occurs between HBr molecules.	0	
	С	Hydrogen bonding is the strongest intermolecular force in liquid ethanol.	0	
	D	Hydrogen bonds occur between C=O and H-N in proteins.	0	
<b>Q2.</b> Th	is que	stion is about intermolecular forces.		(Total 1 mark)
(t	o) E: mo	xplain how permanent dipole-dipole forces arise bet lecules.	ween hydrogen chlo	ride
				(2)

# Q3.

Methanol (CH<sub>3</sub>OH) is an important alcohol with many uses.

(a) Draw a diagram to show how two methanol molecules interact with each other through hydrogen bonding in the liquid phase.

Include all partial charges and all lone pairs of electrons in your diagram.

## Q4.

The ester methyl ethanoate is hydrolysed as shown in the following equation.

 $CH_{3}COOCH_{3}(I) + H_{2}O(I) \iff CH_{3}COOH(I) + CH_{3}OH(I) \qquad \Delta H^{\bullet} = +3 \text{ kJ mol}^{-1}$ 

Which one of the following compounds from the reaction mixture has no hydrogen bonding between its molecules when pure?

0

0

A CH<sub>3</sub>COOCH<sub>3</sub>(I)

**B** H<sub>2</sub>O(I)

C CH<sub>3</sub>COOH(I)

D CH<sub>3</sub>OH(I)

## Q5.

Which molecule does not have a permanent dipole?

- A CH<sub>3</sub>Br
- B CH<sub>2</sub>Br<sub>2</sub>
- C CHBr<sub>3</sub>
- D CBr<sub>4</sub>

(Total 1 mark)

ç0.

(Total 1 mark)

(2)

# Q6.

(a) Methanol has the structure



Explain why the O-H bond in a methanol molecule is polar.

(b) The boiling point of methanol is +65  $^{\circ}$ C; the boiling point of oxygen is -183  $^{\circ}$ C.

Q7.



(b) Crystals of sodium chloride and of diamond both have giant structures. Their melting points are 1074 K and 3827 K, respectively. State the type of structure present in each case and explain why the melting point of diamond is so high.

## Q8.

This question is about intermolecular forces.

 (a) Complete the diagram to show how one molecule of ammonia can form a hydrogen bond with one molecule of ethanol.
 Include all lone pairs of electrons and partial charges on atoms involved in the hydrogen bond.



The table below shows the electronegativity values of atoms of some elements.

Atom	Ĥ	С	Ν	0	Br
Electronegativity	2.1	2.5	3.0	3.5	2.8

- (c) Deduce the **two** atoms from the table above that will form the most polar bond.
- (d) The C–Br bond is polar.

Explain why CBr<sub>4</sub> is **not** a polar molecule.

(e) Suggest, in terms of the intermolecular forces for each compound, why CBr<sub>4</sub> has a

(2)

(1)

(3)

higher boiling point than CHBr<sub>3</sub>



(Total 10 marks)

(2)

## Q9.

Chloroethene can be polymerised to form poly(chloroethene), commonly known as PVC. This polymer can be used to make pipes, window frames and electrical insulation. Plasticisers can be added to change the properties of PVC

A section of poly(chloroethene) is shown.



(a) Chloroethene has a melting point of −154 °C

All types of PVC melt at temperatures over 100 °C

Explain why PVC melts at a higher temperature than chloroethene.

## Q10.

Fritz Haber, a German chemist, first manufactured ammonia in 1909.

Ammonia is very soluble in water.

- (a) State the strongest type of intermolecular force between one molecule of ammonia and one molecule of water.
- (b) Draw a diagram to show how one molecule of ammonia is attracted to one molecule of water. Include all partial charges and all lone pairs of electrons in your diagram.

(c) Phosphine  $(PH_3)$  has a structure similar to ammonia.

In terms of intermolecular forces, suggest the main reason why phosphine is almost insoluble in water.

F.O.

(1) (Total 5 marks)

# Q11.

The table below shows the electronegativity values of some elements.

	Fluorine	Chlorine	Bromine	lodine	Carbon	Hydrogen
Electronegativity	4.0	3.0	2.8	2.5	2.5	2.1

(b) The table below shows the boiling points of fluorine, fluoromethane ( $CH_3F$ ) and hydrogen fluoride.

	F–F	F C H/H H	H–F
Boiling point/K	85	194	293

(i) Name the strongest type of intermolecular force present in:

Liquid F2\_\_\_\_\_\_ Liquid CH3F\_\_\_\_\_\_ Liquid HF\_\_\_\_\_\_ (ii) Explain how the strongest type of intermolecular force in liquid HF arises.

(6)

(c) The table below shows the boiling points of some other hydrogen halides.

	HCI	HBr	Н
Boiling point / K	188	206	238

(i) Explain the trend in the boiling points of the hydrogen halides from HCl to HI.

(ii) Give **one** reason why the boiling point of HF is higher than that of all the other hydrogen halides.

#### Q12

(3) (Total 11 marks)

Van der Waals' forces exist between all molecules. Explain how these forces arise?

(3)

## Q13

This question is about intermolecular forces in some organic compounds.

The table below gives some information about three organic compounds.

Compound	dichloromethane	dichloromethane tetrachloromethane	
Boiling point / °C	40	77	97
Polarity of molecules	polar	non-polar	polar

(c) Explain why tetrachloromethane has a higher boiling point than dichloromethane.

(d) Propan-1-ol has a higher boiling point than the other two compounds because of hydrogen bonding.

Describe the hydrogen bonding in propan-1-ol.



#### 14

The table below shows some values of melting points and some heat energies needed for melting.

Substance	l <sub>2</sub>	NaCl	HF	HCI	НІ
Melting point/K	387	1074	190	158	222
Heat energy for melting /kJ mol-1	7.9	28.9	3.9	2.0	2.9

## (a) Name three types of intermolecular force.

Force 1	
Force 2	
Force 3	

- (b) (i) Describe the bonding in a crystal of iodine.
  - (ii) Name the crystal type which describes an iodine crystal.
  - (iii) Explain why heat energy is required to melt an iodine crystal.

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(3)

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(c) In terms of the intermolecular forces involved, suggest why

(i) hydrogen fluoride requires more heat energy for melting than does hydrogen chloride, (ii) hydrogen iodide requires more heat energy for melting than does hydrogen chloride. (5) Explain why the heat energy required to melt sodium chloride is large. (d) (i) The heat energy needed to vaporise one mole of sodium chloride (171 kJ (ii) mol<sup>-1</sup>) is much greater than the heat energy required to melt one mole of sodium chloride. Explain why this is so. (3) (e) In terms of its structure and bonding, suggest why graphite has a very high melting point. (2) (Total 17 marks)

**15 (b)** Perfume is a mixture of fragrant compounds dissolved in a volatile solvent.

When applied to the skin the solvent evaporates, causing the skin to cool for a short time. After a while, the fragrance may be detected some distance away. Explain these observations.

This	question is about pentan-2-ol and pent-1-ene.	
(a)	The boiling point of pentan-2-ol is 119 °C The boiling point of pent-1-ene is 30 °C	
	Explain why pentan-2-ol has a higher boiling point than pent-1-ene.	
	S	

# Mark schemes

(a)

Power of an atom to attract a pair of electrons in a covalent bond. Allow power of an atom to attract a bonding/shared pair of electrons Allow power of an atom to withdraw electron density from a covalent bond Not lone pair Not Element

(b) <u>Difference in electronegativity</u> leads to bond polarity If chloride (ions) mentioned then CE = 0

(dipoles don't cancel therefore the molecule has an overall permanent dipole) and there is an attraction between  $\partial$ + on one molecule and  $\partial$ - on another

partial charges should be correct if shown and can score M2 from diagram

(c)

SiH <sub>4</sub>	Tetrahedral		1 shape & no tick
PH₃	Pyramidal (trigonal) Allow tetrahedral	$\checkmark$	1 shape & tick
BeCl <sub>2</sub>	Linear		1 shape & no tick
CH₃CI	(Distorted)Tetrahedral	$\checkmark$	1 shape & tick

If shapes are drawn rather than named then penalise first mark gained

[7]

4

[1]

1

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Q3.

(a)

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**M1** on at least one O atom two lone pairs and on at least one OH  $\delta$ + on H and  $\delta$ - on O

**M2** dotted line shown between lone pair on one molecule and the correct H on another

M3 O·····H–O in straight line

Accept pair of dots or crosses for lone pair in place of orbital shape (orbital shape may or may not include two electrons)

Ignore any partial charges on C–H or C–O bonds For straight line in **M3**, allow a deviation of up to 15°

If a different molecule containing hydrogen bonding due to O–H bond drawn (e.g. ethanol, water) or an incorrect attempt at the structure of methanol, then maximum of 2 marks (i.e. only penalise if would score all three marks otherwise)

(b) Idea that lone pairs have greater repulsion than bonding pairs

There must be a comparison between the repulsion of a lone pair and bonding pair

Allow covalent bond = bonding pair

1

1

1

1

(C)

This question is marked using levels of response. Refer to the Mark Scheme Instructions for Examiners for guidance on how to mark this question.				
Level 3 5-6 marks	All stages are covered and the explanation of each stage is generally correct and virtually complete. (6 v 5) Answer is well structured, with no repetition or irrelevant points, and covers all aspects of the question. Accurate and clear expression of ideas with no errors in use of technical terms.			
Level 2 3-4 marks	All stages are covered but stage(s) may be incomplete or may contain inaccuracies OR two			

	stages are covered and are generally correct and virtually complete
	(4 v 3) Answer has some structure and covers most aspects of the question. Ideas are expressed with reasonable clarity with, perhaps, some repetition or some irrelevant points. If any, only minor errors in use of technical terms.
Level 1 1-2 marks	Two stages are covered but stage(s) may be incomplete or may contain inaccuracies OR only one stage is covered but is generally correct and virtually complete
	(2 v 1) Answer includes statements which are presented in a logical order and/or linked.
0 marks	Insufficient correct chemistry to gain a mark.

## Stage 1

Describes the effect of catalyst use

- 1a use of a catalyst has no impact on equilibrium yield
- 1b use of a catalyst gives faster rate
- 1c use of catalyst lowers costs

## Stage 2

Describes the effect of pressure 2a higher pressure gives a higher equilibrium yield 2b higher pressure gives a faster rate 2c the higher the pressure, the greater the cost

## Stage 3

Describes the effect of temperature 3a lower temperature gives a higher equilibrium yield 3b higher temperature gives a faster rate 3c the higher the temperature, the greater the cost Note that converse statements are fine (e.g. 1a higher temperature gives a lower equilibrium yield)

**Q4.** A

[1]

[10]

6

Q5. D

[1]

## Q6.

 Oxygen more/very/highly electronegative (than hydrogen) OR oxygen has stronger attraction for <u>bonding</u> electrons / <u>bonding</u> electrons drawn towards oxygen; causes higher e<sup>-</sup> density round oxygen atom / causes  $H^{+}$  O<sup>\*-</sup>;

(b) van der Waals' forces between oxygen molecules;

Hydrogen bonding between methanol molecules;

H-B stronger than van der Waals' *OR* stronger IMF in methanol; (*if dipole-dipole forces in O*<sub>2</sub> *or methanol, allow comparison, hence max 2*) (*if ionic/covalent etc. max 1*) (*mention of bond break* = CE = 0)

## Q7.

(a) <u>Hydrogen</u> bonding *(full name)* 

Diagram shows at least one  $^{\delta+}H$  and at least one  $^{\delta-}F$ (If full charges shown, M2 = 0)

3 lone pairs shown on at least one fluorine atom H-bond indicated, between H and a lone pair on F

NaCl is ionic (lattice)

(b)

(If atoms not identified, zero for diag) ('FI' for fluorine - mark to Max 2) (Max 1 if only one HF molecule shown, **or** HCI shown)

Dipole results from electronegativity <u>difference</u> **or** values quoted ('difference' may be inferred) (Allow explanation – e.g. F attracts <u>bonding electrons</u> more strongly than H)

**QoL** Fluorine more/very electronegative **or** iodine less electronegative **or** electronegativity difference too small in HI **Comparison required, may be implied**.

HI dipole weaker or bonding e- more equally shared - wtte

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[5]

(Treat atoms/molecules as a contradiction) (Accept 'cubic lattice')
1
Diamond is macromolecular/giant covalent/giant atomic/giant molecular (NOT molecular or tetrahedral) (Ionic/van der Waals' = CE = 0)
1
(Many) covalent/C-C bonds need to be broken / overcome (NOT just 'weakened' etc.) ('Covalent' may be inferred from diagram) (Treat diagram of graphite (without one of diamond) as a contradiction – lose M2 but allow M3/M4])

Which takes much energy **or** covalent bonds are strong (*References to van Der Waals' bonds breaking lose M3/M4*)

[11]

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# Q8.

(a)



M1 – lone pairs and partial charges ( $\delta$ -,  $\delta$ +,  $\delta$ -) on atoms involved in the hydrogen bond

M2 – dotted line between lone pair on N/O to correct H

M3 - linear O-H····N / linear N-H···O

Ignore partial charges on C-H

(b) The (relative) tendency of an atom to attract a pair of electrons/ the electrons/ electron density in a covalent bond

Allow Nucleus instead of atom Power of an atom to attract a bonding/shared pair of electrons Power of an atom to withdraw electron density from a covalent bond

Not lone pair / element

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(c)	H ar	nd O O-H		
(d)	N/1 +	the molecule is completely symmetrical / the molecule is tetrahedral /	1	
(u)	ther	e is an even distribution of electron density	1	
	M2 1	M2 the dipoles cancel out		
		Do not allow	-	
		The polar bonds cancel out / no dipole moment / partial charges cancel		
(e)	M1	CBr <sub>4</sub> has van der Waals' forces between molecules	1	
			1	
	M2	CHBr <sub>3</sub> has van der Waals' forces and dipole-dipole intermolecular forces		
			1	
	М3	The van der Waals' between CBr <sub>4</sub> molecules are stronger than the dipole-dipole and van der Waals' forces between CHBr <sub>3</sub> (because it has a larger mass/more electrons/larger electron cloud)		
		The intermolecular forces between CBr <sub>4</sub> molecules are stronger than the intermolecular forces between CHBr <sub>3</sub>		
		M3 cannot be awarded if mention of breaking bonds		
			1	[10]
				ניטן
Q9.				
(a)	M1	it / PVC is bigger/longer molecule / has more electrons / has bigger surface area / greater <i>M</i> r	1	
			1	
	M2	it / PVC has stronger (van der Waals' / dipole-dipole) forces between		

1

M1 and M2 independent of each other

molecules / intermolecular forces

CE = 0 if reference to hydrogen bonds or breaking of covalent bonds when substances are melted

Comparison must be implied in M1 or M2 to score 2 marks

If there is no comparison at all, then 1 mark could score either for explaining that PVC has strong <u>intermolecular</u> forces due to being a big/long molecule / having many electrons / large surface area / large  $M_r$ , or, for explaining that chloroethene has weak <u>intermolecular</u> forces due to being a small/short molecule / having few electrons / low surface area / low  $M_r$ ,

## ignore additional words

- (c) Need both ideas that
  - it is present AND
  - because PVC needs to be flexible / bendy
     penalise incorrect properties
- (d) Displayed structure required



1

1

1

1

[5]

# Q10.

- (a) Hydrogen bonding / hydrogen bonds / H-bonding / H-Bonds Not just hydrogen.
- (b)



One mark for minimum of 4 correct partial charges shown on the N-H and O-H

One mark for the 3 lone pairs.

One mark for H bond from the lone pair on O or N to the H<sup>5+</sup>

OR



The N-H-O should be linear but can accept if the lone pair on O or N hydrogen bonded to the H If wrong molecules or wrong formula, CE = 0/3

3

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1

1

1

[5]

(c) (Phosphine) does not form hydrogen bonds (with water)

## Q11.

 tendency / strength / ability / power of an <u>atom</u> / <u>element</u> / <u>nucleus</u> to attract / pull / withdraw electrons / e - density / bonding pair / shared pair

in a covalent bond

(b) (i) F<sub>2</sub> = van der Waals' / induced/temporary dipole-dipole / dispersion / London forces

CH<sub>3</sub>F dipole-dipole (not just 'dipole')

HF = hydrogen bonding (not just 'H' / 'hydrogen')

- (ii) large difference in electronegativity between H and F / F most/very/much more electronegative / values '4' & '2.1' quoted (not just 'high<u>er</u>')
  - δ+H-Fδ- dipole created or dipole clearly implied
     (accept arguments such as 'uneven charge in bond'/
     'polar bond' F slightly negative / H slightly positive)

attraction/bond formed between  $\delta$ +H and lone pair on F

		(M2 / M3 may be scored from a diagram)		
		(CE if full charges shown - lose M2 and M3)	1	
(c)	(i)	van der Waals' / induced/temporary dipole-dipole / dispersion / London forces / attractions		
		(ignore references to dipole-dipole)	1	
		increase with the increasing M, / size / mass / Nº of e <sup>-</sup> / size of e <sup>-</sup> cloud (in the hydrogen halides)		
		(if ionic, or if 'covalent bonds broken' = $CE = 0$ ) (mark M1 and M2 separately)		
		(mark wit and wiz separately)	1	
1	(ii)	hydrogen bonding stronger than van der Waals' attraction/forces (accept hydrogen bonding is very strong / strongest) (accept arguments such as 'HF has H-bonds, others <u>only</u> have van der Waals')		
		(not just 'HF has H-bonding')	1	
				[11]

#### 12

- M1 Electron movement in first molecule / temporary dipole
- M2 Induces a dipole in another molecule
- M3 (induced-temporary) attraction of delta + attracts delta in different/adjacent molecules
- M3 depends on M1 and M2
- Allow electrostatic attraction.
- M3 could be scored in diagram.

#### 13

- (c) M1 van der Waals' forces between molecules in CCl<sub>4</sub> stronger than (combined van der Waals' and) dipole-dipole forces between molecules in CH<sub>2</sub>Cl<sub>2</sub>
  - M2 as CCl<sub>4</sub> has (many) more electrons than CH<sub>2</sub>Cl<sub>2</sub>
     M1 must refer to the forces being between molecules at some point
     NOT M1 for any reference to bond breaking
     NOT M1 for any reference to incorrect intermolecular forces
     Allow London forces or temporary (induced) dipole-dipole forces for van der Waals' forces
     For M2, allow CCl<sub>4</sub> has higher mass or higher M<sub>r</sub> or bigger than CH<sub>2</sub>Cl<sub>2</sub>

2

(d) M1 attraction between O lone pair

M2 and δ+ H of OH on another molecule
 no marks if answer indicates that the hydrogen bond is the
 O-H bond within a molecule
 Marks could be awarded from a suitable diagram

#### 14

(a) Force 1: Van der Waals' (1)

Force 2: dipole - dipole (1)

Force 3: hydrogen bonding (1) OR London, Dispersion, temporary dipole

(b) (i) covalent <u>between atoms</u> (1) OR within molecule

Van der Waals' between molecules (1)

- (ii) molecular (1)
- (iii) Bonds (or forces) between molecules must be broken or loosened (1)
   OR V.dW forces
   OR intermolecular forces
   Mention of ions CE=0

(c) (i) H-Bonding in HF (1)

(dipole-) dipole in HCI (1) OR V.dW

H-bonding is stronger than dipole-dipole or V.dW (1) OR H-bonding is a strongest intermolecular force for 3<sup>rd</sup> mark

(ii) HI bigger molecule than HCl **(1)** OR Heavier, more e's, more electron shells, bigger M<sub>r</sub>, more polarisable

Therefore the forces between HI molecules are stronger (1) QL mark (Look for unambiguous statements using correct terminology)

5

2

3

4

(d) (i) ionic **(1)** 

Strong forces between ions (1) OR lots of energy required to break bonds

(ii) All bonds must be broken (1) mention of molecules etc CE=0

3

(e) macromolecular (1)

OR giant molecule / lattice or correct diagram

Strong covalent bonds **(1)** OR lots of energy required to break bonds

## 15

2

4

1

1

1

(b) <u>Solvent</u> has low bp or <u>weak</u> intermolecular forces or evaporates quickly (1)

(Solvent) needs energy <u>to evaporate</u> (to overcome intermolecular forces) or valid reference to latent heat of vaporisation (or evaporation is endothermic) (1)

> OR higher energy or faster molecules more likely to escape so mean energy (and hence temperature) falls

Energy taken from the skin (and so it cools) (1)

Fragrance or perfume (molecules) slowly spreads (through the room) (1)

By random movement or diffusion (of the perfume / fragrance) (1)

#### 16

- (a) M1 idea that pentan-2-ol has stronger intermolecular forces
   M1 idea that hydrogen bonds are stronger than van der Waals' forces
   Penalise M1 for any reference to idea of breaking covalent bonds
  - M2 pent-1-ene has van der Waals' forces (only)
     M2 allow London forces or temporary/induced dipole forces or vdW forces for van der Waals' forces
  - M3 pentan-2-ol (also) has hydrogen bonds M3 Ignore reference to dipole-dipole forces in pentan-2-ol