

# The Advanced Placement Examination in Chemistry

## Part II - Free Response Questions & Answers 1970 to 2006

## Bonding & Molecular Structure

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# Bonding and Molecular Structure

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1973 D

Discuss briefly the relationship between the dipole moment of a molecule and the polar character of the bonds within it. With this as the basis, account for the difference between the dipole moments of  $\text{CH}_2\text{F}_2$  and  $\text{CF}_4$ .

Answer:

In order to have a dipole moment (i.e., to be a polar molecule) a molecule must have polar bonds and must have a molecular geometry which is not symmetrical (i.e., one in which the vector sum of the bond dipoles  $\neq 0$ ).

In  $\text{CH}_2\text{F}_2$  the C-F and C-H bonds are polar and the molecule is not symmetrical; therefore, the molecule is polar and would show a dipole moment.

In  $\text{CF}_4$  the C-F bonds are polar, but the molecule is symmetrical; therefore, the molecule is non-polar and would not show a dipole moment.

1974 D

The possible structures for the compound dinitrogen oxide are NNO and NON. By experimentation it has been found that the molecule of dinitrogen oxide has a non-zero dipole moment and that ions of mass 44, 30, 28, 16, and 14 are obtained in the mass spectrometer. Which of the structures is supported by these data? Show how the data are consistent with this structure.

Answer:

The correct structure is NNO. N-N=O; N-O bond di-pole; non-linear structure, non-symmetrical; molecular dipole moment.

Spectral Data (mass of molecular fragments):

$$44 = \text{NNO} \quad 28 = \text{NN} \quad 14 = \text{N} \quad 30 = \text{NO} \quad 16 = \text{O}$$

A fragment of 28 couldn't be made if the structure was NON.

1974 D

The boiling points of the following compounds increase in the order in which they are listed below:



Discuss the theoretical considerations involved and use them to account for this order.

Answer:

$\text{CH}_4$  - weak London dispersion (van der Waals) forces

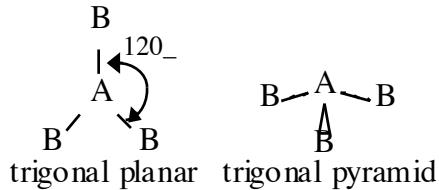
$\text{H}_2\text{S}$  - London forces + dipole-dipole interactions

$\text{NH}_3$  - London + dipole + hydrogen bonding

1975 D

Suppose that a molecule has the formula  $\text{AB}_3$ . Sketch and name two different shapes that this molecule may have. For each of the two shapes, give an example of a known molecule that has that shape. For one of the molecules you have named, interpret the shape in the context of a modern bonding theory.

Answer:



Example: trigonal planar,  $\text{BF}_3$ ; trigonal pyramid,  $\text{NH}_3$

For  $\text{BF}_3$ , the boron atom is surrounded by three pairs of electrons, the arrangement that will minimize the repulsions is a flat (planar) arrangement with the electron pairs furthest apart at  $120^\circ$  angles. **OR**

The  $\text{NH}_3$  molecule has four pairs of electrons: three bonding pairs and one non-bonding pair. The best arrangement for four electron pairs is a tetrahedral structure ( $109.5^\circ$ ) with the lone (non-bonding) electron pair at the apex requiring more space than the bonding pairs, compressing the bonding pairs to an angle of  $107^\circ$ . The

# Bonding and Molecular Structure

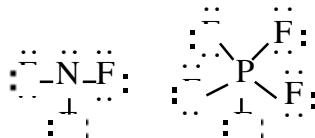
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molecular structure is always based on the positions of the atoms, therefore it is a trigonal pyramid rather than a tetrahedron.

1976 D

$\text{NF}_3$  and  $\text{PF}_5$  are stable molecules. Write the electron-dot formulas for these molecules. On the basis of structural and bonding considerations, account for the fact that  $\text{NF}_3$  and  $\text{PF}_5$  are stable molecules but  $\text{NF}_5$  does not exist.

Answer:



Describe the  $sp^3$  bonding for  $\text{NF}_3$  and the  $sp^3d$  for  $\text{PF}_5$ .

Nonexistence of  $\text{NF}_5$  because of no low energy  $d$  orbital for N.

1978 D

State precisely what is meant by each of the following four terms. Then distinguish clearly between each of the two terms in part (a) and between each of the two terms in part (b), using chemical equations or examples where helpful.

- Bond polarity and molecular polarity (dipole moment)
- For a metal M, ionization energy and electrode potential.

Answer:

(a) Bond polarity - resulting from unequal sharing of electrons between bonding atoms; or from bonding of atoms with different electronegativities.

Molecular polarity - result of the separation of the centers of positive and negative charges in an entire molecule (The dipole moment is a measure); or the result of the non-zero vector sum of bond dipoles and lone-pair electrons.

Distinction (normally included within the definitions).

(b) Ionization energy - energy required to remove an electron from an atom [if atom is described as gaseous or isolated - 1 additional point]

Electrode potential - related to energy associated with oxidation or reduction or associated with a tendency to gain or lose electrons.

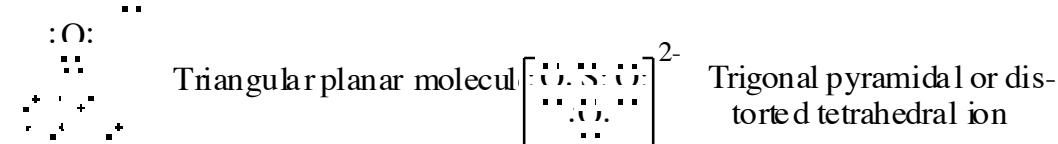
A quantity measured relative to the hydrogen electrode or related to the energy changes in an electrochemical cell.

Distinction (normally included within the definitions).

1979 D

Draw Lewis structures for  $\text{CO}_2$ ,  $\text{H}_2$ ,  $\text{SO}_3$  and  $\text{SO}_3^{2-}$  and predict the shape of each species.

Answer:



1979 D

# Bonding and Molecular Structure

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Butane, chloroethane, acetone, and 1-propanol all have approximately the same molecular weights. Data on their boiling points and solubilities in water are listed in the table below.

Compound	Formula	Boiling Pt.(°C)	Solubility in water
Butane	CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>	0	insoluble
Chloroethane	CH <sub>3</sub> CH <sub>2</sub> Cl	12	insoluble
Acetone	$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3\text{C}=\text{C}\text{CH}_3 \end{array}$	56	completely miscible
1-Propanol	CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> OH	97	completely miscible

On the basis of dipole moments (molecular polarities) and/or hydrogen bonding, explain in a qualitative way the differences in the

- (a) boiling points of butane and chloroethane.
- (b) water solubilities of chloroethane and acetone.
- (c) water solubilities of butane and 1-propanol.
- (d) boiling points of acetone and 1-propanol.

Answer:

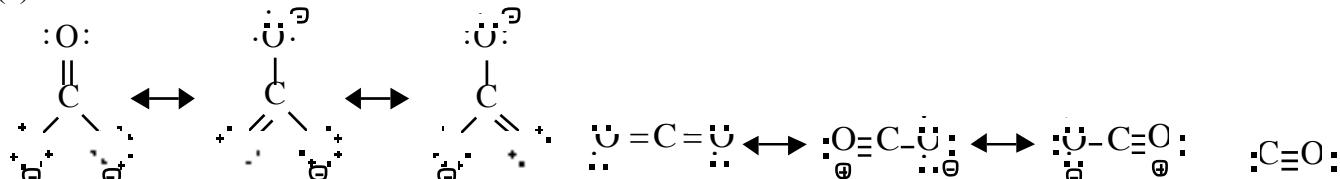
- (a) Butane is nonpolar; chloroethane is polar. Intermolecular forces of attraction in liquid chloroethane are larger due to dipole-dipole attraction; thus a higher boiling point for chloroethane.
- (b) Both chloroethane and acetone are polar. However, acetone forms hydrogen bonds to water much more effectively than chloroethane does, resulting in greater solubility of acetone in water.
- (c) Butane is non-polar and cannot form hydrogen bonds; 1-propanol is polar and can form hydrogen bonds. 1-propanol can interact with water by both dipole-dipole forces and hydrogen bonds. Butane can interact with water by neither means. Thus, 1-propanol is much more soluble.
- (d) Acetone molecules are attracted to each other by van der Waals attraction and dipole-dipole attraction. 1-propanol molecules show these two types of attraction. However, 1-propanol can also undergo hydrogen bonding. This distinguishing feature results in the higher boiling point of 1-propanol.

1982 D

- (a) Draw the Lewis electron-dot structures for CO<sub>3</sub><sup>2-</sup>, CO<sub>2</sub>, and CO, including resonance structures where appropriate.
- (b) Which of the three species has the shortest C-O bond length? Explain the reason for your answer.
- (c) Predict the molecular shapes for the three species. Explain how you arrived at your predictions.

Answer:

- (a)



- (b) CO has the shortest bond because there is a triple bond. OR because there is the greatest number of electrons between C and O in CO.
- (c) CO<sub>3</sub><sup>2-</sup> trigonal planar (planar and triangular). C bonding is *sp*<sup>2</sup> hybrid - or - C has three bonding pairs and no lone pair.

CO<sub>2</sub> linear. C bonding is *sp* hybrid - or - C has two bonding pairs and no lone pairs - or - CO<sub>2</sub> is nonpolar and must be linear.

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CO linear. Two atoms determine a straight line.

1982 D

The values of the first three ionization energies ( $I_1$ ,  $I_2$ ,  $I_3$ ) for magnesium and argon are as follows:

	$I_1$ (kJ/mol)	$I_2$ (kJ/mol)	$I_3$ (kJ/mol)
Mg	735	1443	7730
Ar	1525	2665	3945

- (a) Give the electronic configurations of Mg and Ar.
- (b) In terms of these configurations, explain why the values of the first and second ionization energies of Mg are significantly lower than the values for Ar, whereas the third ionization energy of Mg is much larger than the third ionization energy of Ar.
- (c) If a sample of Ar in one container and a sample of Mg in another container are each heated and chlorine is passed into each container, what compounds, if any, will be formed? Explain in terms of the electronic configurations given in part (a).
- (d) Element Q has the following first three ionization energies:

	$I_1$ (kJ/mol)	$I_2$ (kJ/mol)	$I_3$ (kJ/mol)
Q	496	4568	6920

What is the formula for the most likely compound of element Q with chlorine? Explain the choice of formula on the basis of the ionization energies.

Answer:

- (a) Mg:  $1s^2 2s^2 2p^6 3s^1$   
Ar:  $1s^2 2s^2 2p^6 3s^2 3p^6$
- (b) Valence electrons for Mg and Ar are in the same principal energy level, but Ar atom is smaller and has a greater nuclear charge. Thus, ionization energies for Mg are less than those for Ar. Removal of third electron from Mg atom is from  $n = 2$  level and electrons in this level experience strong nuclear attraction.
- (c) Only  $MgCl_2$  forms. Mg atoms readily lose 2 valence electrons each. Ionization energy for third electron very high. Electron affinity for Ar is low, and ionization energies for Ar atoms are high.
- (d) Formula is  $QCl$ . Very high second ionization energy indicates that there is only one valence electron.

1985 D

Substance	Melting Point, °C
$H_2$	-259
$C_3H_8$	-190
HF	-92
CsI	621
LiF	870
SiC	>2,000

- (a) Discuss how the trend in the melting points of the substances tabulated above can be explained in terms of the types of attractive forces and/or bonds in these substances.
- (b) For any pairs of substances that have the same kind(s) of attractive forces and/or bonds, discuss the factors that cause variations in the strengths of the forces and/or bonds.

Answer:

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- (a) H<sub>2</sub> and C<sub>3</sub>H<sub>8</sub> have low melting points because the forces involved were the weak van der Waals (or London) forces.

HF has a higher melting point because intermolecular hydrogen bonding is important.

CsI and LiF have still higher melting points because ionic lattice forces must be overcome to break up the crystals, and the ionic forces are stronger than van der Waals forces and hydrogen bonds.

SiC is an example of a macromolecular substance where each atom is held to its neighbors by very strong covalent bonds.

- (b) C<sub>3</sub>H<sub>8</sub> and H<sub>2</sub>: There are more interactions per molecule in C<sub>3</sub>H<sub>8</sub> than in H<sub>2</sub>. OR C<sub>3</sub>H<sub>8</sub> is weakly polar and H<sub>2</sub> is nonpolar.

LiF and CsI: The smaller ions in LiF result in a higher lattice energy than CsI has. Lattice energy U is proportional to  $\frac{1}{r^+ + r^-}$ .

1988 D

Using principles of chemical bonding and/or intermolecular forces, explain each of the following.

- (a) Xenon has a higher boiling point than neon has.

- (b) Solid copper is an excellent conductor of electricity, but solid copper chloride is not.

- (c) SiO<sub>2</sub> melts at a very high temperature, while CO<sub>2</sub> is a gas at room temperature, even though Si and C are in the same chemical family.

- (d) Molecules of NF<sub>3</sub> are polar, but those of BF<sub>3</sub> are not.

Answer:

- (a) Xe and Ne are monatomic elements held together by London dispersion (van der Waals) forces. The magnitude of such forces is determined by the number of electrons in the atom. A Xe atom has more electrons than a neon atom has. (Size of the atom was accepted but mass was not.)

- (b) The electrical conductivity of copper metal is based on mobile valence electrons (partially filled bands). Copper chloride is a rigid ionic solid with the valence electrons of copper localized in individual copper(II) ions.

- (c) SiO<sub>2</sub> is a covalent network solid. There are strong bonds, many of which must be broken simultaneously to volatilize SiO<sub>2</sub>. CO<sub>2</sub> is composed of discrete, nonpolar CO<sub>2</sub> molecules so that the only forces holding the molecules together are the weak London dispersion (van der Waals) forces.

- (d) In NF<sub>3</sub>, a lone pair of electrons on the central atom results in a pyramidal shape. The dipoles don't cancel, thus the molecule is polar.

While in BF<sub>3</sub>, there is no lone pair on the central atom so the molecule has a trigonal planar shape in which the dipoles cancel, thus the molecule is nonpolar.

1989 D



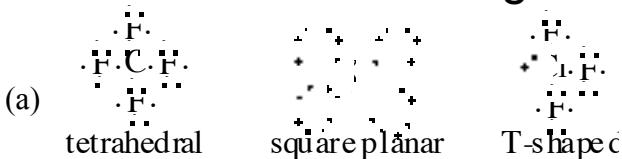
- (a) Draw a Lewis electron-dot structure for each of the molecules above and identify the shape of each.

- (b) Use the valence shell electron-pair repulsion (VSEPR) model to explain the geometry of each of these molecules.

Answer:

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- (b)  $\text{CF}_4$  = 4 bonding pairs around C at corners of regular tetrahedron to minimize repulsion (maximize bond angles).  
 $\text{XeF}_4$  = 4 bonding pairs and 2 lone pairs give octahedral shape with lone pairs on opposite sides of Xe atom  
 $\text{ClF}_3$  = 3 bonding pairs and 2 lone pairs give trigonal bipyramidal with one pairs in equatorial positions  $120^\circ$  apart.

1989 D

The melting points of the alkali metals decrease from Li to Cs. In contrast, the melting points of the halogens increase from  $\text{F}_2$  to  $\text{I}_2$ .

- (a) Using bonding principles, account for the decrease in the melting points of the alkali metals.  
(b) Using bonding principles, account for the decrease in the melting points of the halogens.  
(c) What is the expected trend in the melting points of the compounds  $\text{LiF}$ ,  $\text{NaCl}$ ,  $\text{KBr}$ , and  $\text{CsI}$ ? Explain this trend using bonding principles.

Answer:

- (a) Alkali metals have metallic bonds: cations in a sea of electrons. As cations increase in size (Li to Cs), charge density decreases and attractive forces (and melting points) decrease.  
(b) Halogen molecules are held in place by dispersion (van der Waals) forces: bonds due to temporary dipoles caused by polarization of electron clouds. As molecules increase in size ( $\text{F}_2$  to  $\text{I}_2$ ), the larger electron clouds are more readily polarized, and the attractive forces (and melting points) increase.  
(c) Melting point order:  $\text{LiF} > \text{NaCl} > \text{KBr} > \text{CsI}$   
Compounds are ionic. Larger radii of ions as listed. Larger radii  $\rightarrow$  smaller attraction and lower melting points.

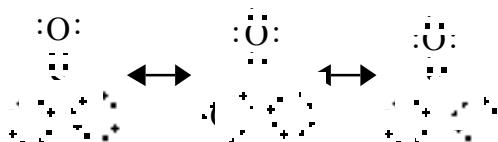
1990 D (Required)

Use simple structure and bonding models to account for each of the following.

- (a) The bond length between the two carbon atoms is shorter in  $\text{C}_2\text{H}_4$  than in  $\text{C}_2\text{H}_6$ .  
(b) The H-N-H bond angle is  $107.5^\circ$ , in  $\text{NH}_3$ .  
(c) The bond lengths in  $\text{SO}_3$  are all identical and are shorter than a sulfur-oxygen single bond.  
(d) The  $\text{I}_3^-$  ion is linear.

Answer:

- (a)  $\text{C}_2\text{H}_4$  has a multiple bond;  $\text{C}_2\text{H}_6$  has a single bond. Multiple bonds are stronger and, therefore, shorter than single bonds.  
(b)  $\text{NH}_3$  has 3 bonding pairs of electrons and 1 lone pair. Bonding pairs are forced together because repulsion between lone pair and bonding pairs is greater than between bonding pairs.  
(c) The bonding in  $\text{SO}_3$  can be described as a combination of 3 resonance forms of 1 double and 2 single bonds.



## Bonding and Molecular Structure

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The actual structure is intermediate among the 3 resonance forms, having 3 bonds that are equal and stronger (therefore, shorter) than an S-O single bond.

- (d) The central I atom has 3 lone pairs and 2 bonding pairs around it. : $\ddot{\text{I}}$ : $\ddot{\text{I}}$ : $\ddot{\text{I}}$ :

To minimize repulsion, the 3 lone pairs on the central atom are arranged as a triangle in a plane are right angles to the I-I-I- axis.

1991 D

Experimental data provide the basis for interpreting differences in properties of substances.

TABLE 1

Compound	Melting Point (°C)	Electrical Conductivity of Molten State (ohm <sup>-1</sup> )
BeCl <sub>2</sub>	405	0.086
MgCl <sub>2</sub>	714	> 20
SiCl <sub>4</sub>	-70	0
MgF <sub>2</sub>	1261	> 20

TABLE 2

Substance	Bond Length (angstroms)
F <sub>2</sub>	1.42
Br <sub>2</sub>	2.28
N <sub>2</sub>	1.09

Account for the differences in properties given in Tables 1 and 2 above in terms of the differences in structure and bonding in each of the following pairs.

- (a) MgCl<sub>2</sub> and SiCl<sub>4</sub>      (c) F<sub>2</sub> and Br<sub>2</sub>  
 (b) MgCl<sub>2</sub> and MgF<sub>2</sub>      (d) F<sub>2</sub> and N<sub>2</sub>

**Answer:**

- (a)  $\text{MgCl}_2$  is IONIC while  $\text{SiCl}_4$  is COVALENT. The electrostatic, interionic forces in magnesium chloride are much stronger than the intermolecular (dispersion) forces in  $\text{SiCl}_4$  and lead to a higher melting point for  $\text{MgCl}_2$ . Molten  $\text{MgCl}_2$  contains mobile ions that conduct electricity whereas molten  $\text{SiCl}_4$  is molecular, not ionic, and has no conductivity.
  - (b)  $\text{MgF}_2$  has a higher melting point than  $\text{MgCl}_2$  because the smaller  $\text{F}^-$  ions and the smaller interionic distances in  $\text{MgF}_2$  cause stronger forces and higher melting point.
  - (c) The bond length in  $\text{Br}_2$  is larger than in  $\text{F}_2$  because the Br atom is bigger (more shells) than the F atom.
  - (d) The bond length in  $\text{N}_2$  is less than in  $\text{F}_2$  because the N≡N bond is triple and the F-F is single. Triple bonds are stronger and therefore shorter than single bonds.

1992 D

Explain each of the following in terms of atomic and molecular structures and/or intermolecular forces.

- (a) Solid K conducts an electric current, whereas solid  $\text{KNO}_3$  does not.
  - (b)  $\text{SbCl}_3$  has measurable dipole moment, whereas  $\text{SbCl}_5$  does not.
  - (c) The normal boiling point of  $\text{CCl}_4$  is  $77^\circ\text{C}$ , whereas that of  $\text{CBr}_4$  is  $190^\circ\text{C}$ .
  - (d)  $\text{NaI(s)}$  is very soluble in water, whereas  $\text{I}_2\text{(s)}$  has a solubility of only 0.03 gram per 100 grams of water.

**Answer:**

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- (a) K conducts because of its metallic bonding - or - “sea” of mobile electrons (or free electrons).  $\text{KNO}_3$  does not conduct because it is ionically bonded and has immobile ions (or immobile electrons).
- (b)  $\text{SbCl}_3$  has a measurable dipole moment because it has a lone pair of electrons which causes a dipole - or - its dipoles do not cancel - or - it has a trigonal pyramidal structure - or - a clear diagram illustrating any of the above.
- (c)  $\text{CBr}_4$  boils at a higher temperature than  $\text{CCl}_4$  because it has stronger intermolecular forces (or van der Waal or dispersion). These stronger forces occur because  $\text{CBr}_4$  is larger and/or has more electrons than  $\text{CCl}_4$ .
- (d)  $\text{NaI}$  has greater aqueous solubility than  $\text{I}_2$  because  $\text{NaI}$  is ionic (or polar), whereas  $\text{I}_2$  is non-polar (or covalent). Water, being polar, interacts with the ions of  $\text{NaI}$  but not with  $\text{I}_2$ . (Like dissolves like accepted if polarity of water is clearly indicated.)

## 1992 D



Nitrogen is the central atom in each of the species given above.

- (a) Draw the Lewis electron-dot structure for each of the three species.
- (b) List the species in order of increasing bond angle. Justify your answer.
- (c) Select one of the species and give the hybridization of the nitrogen atom in it.
- (d) Identify the only one of the species that dimerizes and explain what causes it to do so.

Answer:



(b)  $\text{NO}_2^- < \text{NO}_2 < \text{NO}_2^+$

$\text{NO}_2^-$  - 3 charge centers around N; lone pair of electrons on N

$\text{NO}_2$  - 3 charge centers around N; single electron on N

$\text{NO}_2^+$  - 2 charge centers on N

(c)  $\text{NO}_2^+$  is linear, has  $sp$  hybridization - or -  $\text{NO}_2/\text{NO}_2^-$  have  $sp^2$  hybridization

(d)  $\text{NO}_2$  will dimerize, because it contains an odd electron that will pair readily with another, forming  $\text{N}_2\text{O}_4$ .

## 1994 D

Use principles of atomic structure and/or chemical bonding to answer each of the following.

- (a) The radius of the Ca atom is 0.197 nanometer; the radius of the  $\text{Ca}^{2+}$  ion is 0.099 nanometer. Account for this difference.
- (b) The lattice energy of  $\text{CaO}_{(s)}$  is -3,460 kilojoules per mole; the lattice energy for  $\text{K}_2\text{O}_{(s)}$  is -2,240 kilojoules per mole. Account for this difference.

Ionization Energy (kJ/mol)		
	First	Second
K	419	3,050
Ca	590	1,140

- (c) Explain the difference between Ca and K in regard to
- their first ionization energies,
  - their second ionization energies.
- (d) The first ionization energy of Mg is 738 kilojoules per mole and that of Al is 578 kilojoules per mole. Account for this difference.

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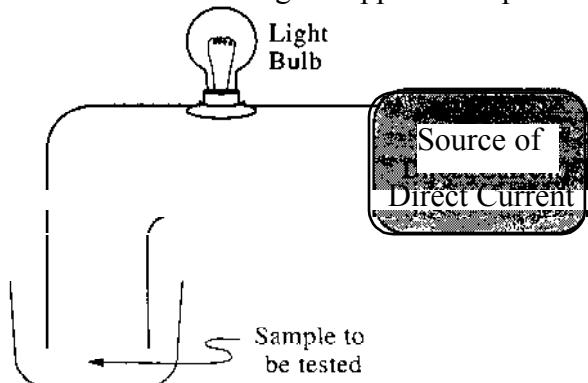
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Answer:

- (a) The valence electrons in a calcium atom are the  $4s^2$ . In a calcium ion these electrons are absent and the highest energy electrons are  $3p$ , which has a much smaller size because the  $(-)/(+)$  charge ratio is less than 1 causing a contraction of the electron shell.
- (b) Lattice energy can be represented by Coulomb's law: lattice energy =  $k \frac{Q_1 Q_2}{r}$ , where  $Q_1$  and  $Q_2$  are the charges on the ions, in CaO these are +2 and -2 respectively, while in K<sub>2</sub>O they are +1 and -2. The  $r$  (the distance between ions) is slightly smaller in CaO, combined with the larger charges, thus accounts for the larger lattice energy.
- (c) Electron arrangements: K = [Ar]  $4s^1$ , Ca = [Ar]  $4s^2$
- (i) Potassium has a single  $4s$  electron that is easily removed to produce an [Ar] core, whereas, calcium has paired  $4s$  electrons which require greater energy to remove one.
- (ii) a K<sup>+</sup> ion has a stable [Ar] electron core and requires a large amount of energy to destabilize it and create a K<sup>2+</sup> ion. Ca<sup>+</sup> has a remaining  $4s^1$  electron that is more easily removed than a core electron, but not as easily as its first  $4s$  electron.
- (d) Electron arrangements,
- Mg = [Ne]  $3s^2$ , Al = [Ne]  $3s^2, 3p^1$
- It is easier to remove a shielded, single, unpaired  $3p$  electron from the aluminum than to remove one electron from a paired  $3s$  orbital in magnesium.

## 1995 D (Required)

The conductivity of several substances was tested using the apparatus represented by the diagram below.



The results of the tests are summarized in the following data table.

	AgNO <sub>3</sub>	Sucrose	Na	H <sub>2</sub> SO <sub>4</sub> (98%)
Melting Point (°C)	212°	185°	99°	Liquid at Room Temp.
Liquid (fused)	++	-	++	+
Water Solution	++	-	++ <sup>(1)</sup>	++ <sup>(2)</sup>
Solid	-	-	++	Not Tested

Key:    ++ Good conductor  
          + Poor conductor  
          - Nonconductor

(1) Dissolves, accompanied by evolution of flammable gas

(2) Conduction increases as the acid is added slowly

# Bonding and Molecular Structure

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and carefully to water

Using models of chemical bonding and atomic or molecular structure, account for the differences in conductivity between the two samples in each of the following pairs.

- (a) Sucrose solution and silver nitrate solution.
- (b) Solid silver nitrate and solid sodium metal.
- (c) Liquid (fused) sucrose and liquid (fused) silver nitrate.
- (d) Liquid (concentrated) sulfuric acid and sulfuric acid solution.

Answer:

- (a) Sucrose, composed of all non-metals, is a covalently bonded molecule that does not ionize in water and, therefore, does not produce a conducting solution. Silver nitrate has an ionic bond between the silver cation and the nitrate anion that is hydrated in water producing a conducting ionic solution.
- (b) Silver nitrate has covalent bonds in the nitrate anion and an ionic bond between the cation and anion but in the solid state these ions are not free to move and conduct an electric current. In sodium, a metal crystal, it has a large number of closely spaced molecular orbitals that form a virtual continuum of levels called bands. Empty molecular orbitals are close in energy to filled molecular orbitals. Mobile electrons are furnished when electrons in filled molecular orbitals are excited into empty ones. These conduction electrons are free to travel throughout the metal crystal
- (c) Fused sucrose does not contain any ions to carry an electrical charge whereas the ions in silver nitrate are now free to move in the liquid and conduct the charge.
- (d) Concentrated sulfuric acid has very little water to hydrolyze its ions and is only slightly ionized. As it is added to water, appreciable amounts of ions are present as the molecular  $\text{H}_2\text{SO}_4$  is dissociated into hydrogen and sulfate ions.

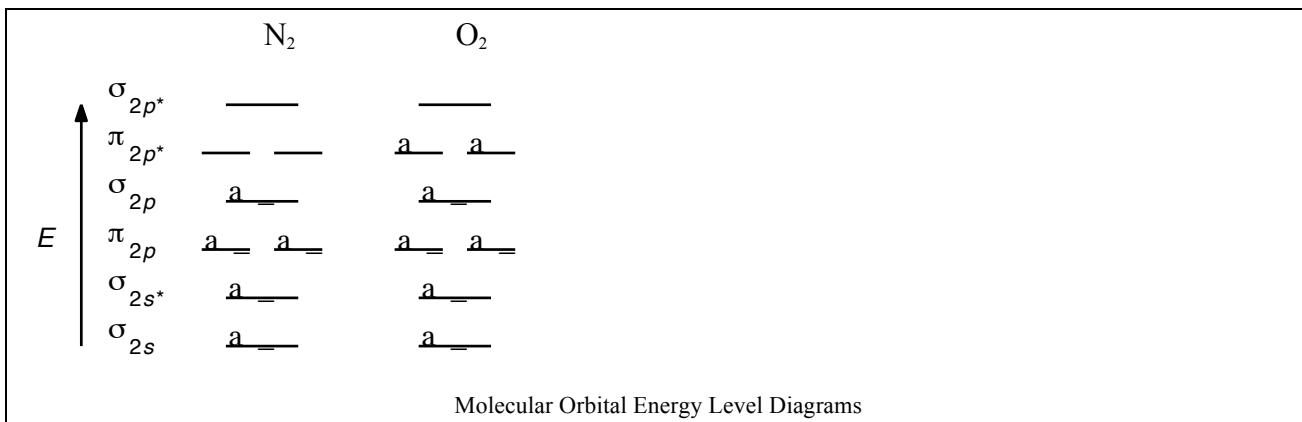
## 1995 D

Explain the following in terms of the electronic structure and bonding of the compounds considered.

- (a) Liquid oxygen is attracted to a strong magnet, whereas liquid nitrogen is not.
- (b) The  $\text{SO}_2$  molecule has a dipole moment, whereas the  $\text{CO}_2$  molecule has no dipole moment. Include the Lewis (electron-dot) structures in your explanation.
- (c) Halides of cobalt(II) are colored, whereas halides of zinc(II) are colorless.
- (d) A crystal of high purity silicon is a poor conductor of electricity; however, the conductivity increases when a small amount of arsenic is incorporated (doped) into the crystal.

Answer:

(a)

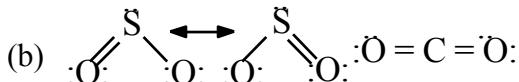


Paramagnetism causes a substance to be attracted into the inducing magnetic field. Paramagnetism is associated with unpaired electrons, as in oxygen but diamagnetism (repelled from the inducing magnetic

# Bonding and Molecular Structure

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field) is associated with paired electrons as in nitrogen. Any substance that has both paired and unpaired electrons will exhibit paramagnetism, since that effect is stronger than diamagnetism.



There is a dipole moment between the oxygen and the sulfur in sulfur dioxide and a bond angle of  $119^\circ$ . This results in a net dipole in the molecule. While there is a dipole in the carbon-oxygen bond, the  $180^\circ$  bond angle cancels the dipole moment in the molecule.

- (c) A cobalt(II) ion has the electron configuration of [Ar]  $3d^7$ . It has 2 paired  $d$  electrons and 3 unpaired electrons. According to crystal field theory, as the chloride ion approaches the cobalt(II) ion, repulsion between the chloride lone pairs and the metal electrons affects the metal  $d$  orbitals differently (the  $x^2-y^2$  and  $z^2$  more than the  $xy$ ,  $xz$ , and  $yz$ ). There is an energy difference between the sets of  $d$  orbitals. The energy difference between sets of  $d$  orbitals is comparable to the energy of visible light. In zinc ions, all the  $d$  orbitals are paired and all the orbitals are degenerate.
- (d) Arsenic atoms have one more valence electron than silicon atoms and can lose an electron to form  $\text{As}^+$  ions which can occupy some of the lattice points in the silicon crystal. If the amount of arsenic is kept small then these ions don't interact. The extra electrons from the arsenic occupy orbitals in a narrow band of energies that lie between the filled and empty bands of the silicon. This structure decreases the amount of energy required to excite an electron into the lowest-energy empty band in the silicon and increases the number of electrons that have enough energy to cross this gap.

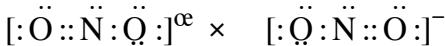
## 1996 D

Explain each of the following observations in terms of the electronic structure and/or bonding of the compounds involved.

- (a) At ordinary conditions, HF (normal boiling point =  $20^\circ\text{C}$ ) is a liquid, whereas HCl (normal boiling point = -  $114^\circ\text{C}$ ) is a gas.
- (b) Molecules of  $\text{AsF}_3$  are polar, whereas molecules of  $\text{AsF}_5$  are nonpolar.
- (c) The N-O bonds in the  $\text{NO}_2^-$  ion are equal in length, whereas they are unequal in  $\text{HNO}_2$ .
- (d) For sulfur, the fluorides  $\text{SF}_2$ ,  $\text{SF}_4$ , and  $\text{SF}_6$  are known to exist, whereas for oxygen only  $\text{OF}_2$  is known to exist.

Answer:

- (a) hydrogen bonding (dipole-dipole attraction) is much larger in HF than in HCl.
- (b)  $\text{AsF}_3$  forms a pyramidal shaped molecule with a lone pair of electrons creating an asymmetrical region opposite the three highly electron-affinitive fluorine in the base. The  $\text{AsF}_5$  molecule has a highly symmetrical trigonal bipyramidal shape with no lone electron pairs.
- (c) The N-O bonds in the nitrite ion are stabilized by resonance and are of equal length, but in  $\text{HNO}_2$ , with a hydrogen attached to an oxygen, resonance is no longer possible.



- (d) There are only four orbitals in the valence shell of oxygen, one  $s$  and three  $p$ 's. As a result, oxygen can hold no more than eight valence electrons, which it gets when it forms  $\text{OF}_2$ . The valence orbitals of sulfur are in the  $n = 3$  shell and includes empty  $d$  orbitals that can be used to expand its valence shell. Sulfur has 10 valence electrons in forming  $\text{SF}_4$  and 12 valence electrons to form  $\text{SF}_6$  ( $sp^3d^2$  hybrid orbitals).

## 1997 D (Required)

Consider the molecules  $\text{PF}_3$  and  $\text{PF}_5$ .

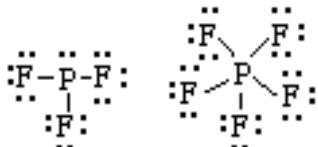
# Bonding and Molecular Structure

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- (a) Draw the Lewis electron-dot structures for  $\text{PF}_3$  and  $\text{PF}_5$  and predict the molecular geometry of each.
- (b) Is the  $\text{PF}_3$  molecule polar, or is it nonpolar? Explain.
- (c) On the basis of bonding principles, predict whether each of the following compounds exists. In each case, explain your prediction.
- $\text{NF}_5$
  - $\text{AsF}_5$

Answer:

- (a)  $\text{PF}_3$  = tripod (pyramid);  $\text{PF}_5$  = trigonal bipyramid



- (b) polar; net dipole moment toward the non-symmetrical position of the fluorines.
- (c) (i)  $\text{NF}_5$  - doesn't exist, nitrogen can't hybridize to form the  $dsp^3$  orbitals and is also too small to accommodate 5 fluorine atoms around it.
- (ii)  $\text{AsF}_5$  - does exist, arsenic can hybridize to form the  $dsp^3$  orbitals and is large enough to accommodate 5 fluorine atoms around it.

## 1998 D

Answer each of the following using appropriate chemical principles.

- (c) Dimethyl ether,  $\text{H}_3\text{C-O-CH}_3$ , is not very soluble in water. Draw a structural isomer of dimethyl ether that is much more soluble in water and explain the basis of its increased water solubility.

In each case, justify your choice.

Answer

- (c)   
The O-H bond in ethyl alcohol is very polar and will allow the molecule to be attracted to and dissolve in the polar water.

## 1999 B

Answer the following questions regarding light and its interactions with molecules, atoms, and ions.

- (a) The longest wavelength of light with enough energy to break the Cl–Cl bond in  $\text{Cl}_2(g)$  is 495 nm.
- Calculate the frequency, in  $\text{s}^{-1}$ , of the light.
  - Calculate the energy, in J, of a photon of the light.
  - Calculate the minimum energy, in  $\text{kJ mol}^{-1}$ , of the Cl–Cl bond.
- (b) A certain line in the spectrum of atomic hydrogen is associated with the electronic transition of the H atom from the sixth energy level ( $n = 6$ ) to the second energy level ( $n = 2$ ).
- Indicate whether the H atom emits energy or whether it absorbs energy during the transition. Justify your answer.
  - Calculate the wavelength, in nm, of the radiation associated with the spectral line.
  - Account for the observation that the amount of energy associated with the same electronic transition ( $n = 6$  to  $n = 2$ ) in the  $\text{He}^+$  ion is greater than that associated with the corresponding transition in the H atom.

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## Answer

(a) (i)  $c = \lambda v ; v = \frac{c}{?} = \frac{3.00 \times 10^8 \text{ m/s}}{495 \times 10^9 \text{ m}} = 6.06 \times 10^{14} \text{ waves/sec}$

(ii)  $\Delta E = \hbar v = (6.63 \times 10^{-34} \text{ joule}\cdot\text{s})(6.06 \times 10^{14} \text{ s}^{-1}) = 4.02 \times 10^{-19} \text{ J}$

(iii)  $(4.02 \times 10^{-19} \text{ J})(6.02 \times 10^{23} \text{ mol}^{-1}) = 241,870 \text{ J} = 242 \text{ kJ}$

- (b) (i) emits energy. The  $n = 6$  state is at a higher energy than  $n = 2$ , the electron must release energy go to the lower state.

(ii)  $E_2 = \frac{2.178 \times 10^{18} \text{ J}}{2^2} = -5.445 \times 10^{-19} \text{ J}$

$E_6 = \frac{2.178 \times 10^{18} \text{ J}}{6^2} = -6.05 \times 10^{-20} \text{ J}$

$\Delta E = E_6 - E_2 = 4.84 \times 10^{-19} \text{ J}$

## OR

$$\Delta E = -2.178 \times 10^{-18} \left( \frac{1}{2^2} - \frac{1}{6^2} \right) \text{ J} = 4.84 \times 10^{-19} \text{ J}$$

## OR

$$\lambda = \frac{hc}{E}; v = \frac{E}{h} = \frac{4.84 \times 10^{-19} \text{ J}}{6.626 \times 10^{-34} \text{ J}\cdot\text{sec}} =$$

$$= 7.30 \times 10^{14} \text{ sec}^{-1}$$

$$\lambda = \frac{c}{v} = \frac{3.00 \times 10^{-18} \text{ m/sec}}{7.30 \times 10^{14} \text{ sec}^{-1}} = 4.11 \times 10^{-7} \text{ m}$$

## OR

$$\lambda = \frac{(6.626 \times 10^{-34} \text{ J}\cdot\text{sec})(3.00 \times 10^{17} \text{ nm/sec})}{4.84 \times 10^{-19} \text{ J}} =$$

$$= 411 \text{ nm}$$

- (iii) the  $\text{He}^+$  has a nuclear charge of 2+ vs H with a 1+, therefore, it has a stronger hold on the electron. This requires more energy for the electron to move to higher energy levels and when the electron moves from higher to lower energies, it releases more energy.

## 1999 D

Answer the following questions using principles of chemical bonding and molecular structure.

- (a) Consider the carbon dioxide molecule,  $\text{CO}_2$ , and the carbonate ion,  $\text{CO}_3^{2-}$ .

(i) Draw the complete Lewis electron-dot structure for each species.

(ii) Account for the fact at the carbon-oxygen bond length in  $\text{CO}_3^{2-}$  is greater than the carbon-oxygen bond length in  $\text{CO}_2$ .

- (b) Consider the molecules  $\text{CF}_4$  and  $\text{SF}_4$ .

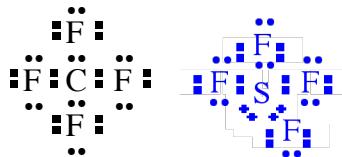
(i) Draw the complete Lewis electron-dot structure for each molecule.

(ii) In terms of molecular geometry, account for the fact that the  $\text{CF}_4$  molecule is nonpolar, whereas the  $\text{SF}_4$  molecule is polar.

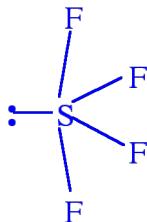
# Bonding and Molecular Structure

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Answer

- (a) (i)  there are two other similar resonance structures for the carbonate ion.
- (ii) the pi O=C double bond in CO<sub>2</sub> is shorter than a single O-C resonance sigma-bond (all are identical and are about 1 1/3 bond) found in a carbonate ion.
- (b) (i) 

(ii) in the tetrahedral CF<sub>4</sub>, the polar C-F bonds are cancelled out by the equiangular pull of the 4 bonds. With an expanded octet and trigonal bipyramidal structure, SF<sub>4</sub> has a pair of unbonded electrons at the center of the bipyramid, this gives a “seasaw” shape to the molecule and an uneven pull to the polar S-F bonds.



## 2000 D

Answer the following questions about the element selenium, Se (atomic number 34).

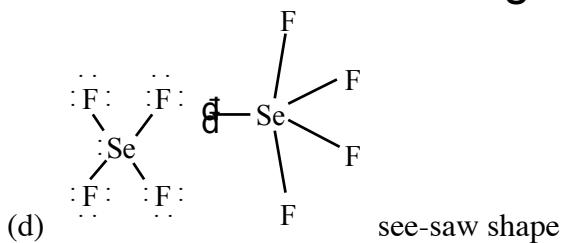
- (a) Samples of natural selenium contain six stable isotopes. In terms of atomic structure, explain what these isotopes have in common, and how they differ.
- (b) Write the complete electron configuration (e.g., 1s<sup>2</sup> 2s<sup>2</sup>... etc.) for a selenium atom in the ground state. Indicate the number of unpaired electrons in the ground-state atom, and explain your reasoning.
- (c) In terms of atomic structure, explain why the first ionization energy of selenium is
- less than that of bromine (atomic number 35), and
  - greater than that of tellurium (atomic number 52).
- (d) Selenium reacts with fluorine to form SeF<sub>4</sub>. Draw the complete Lewis electron-dot structure for SeF<sub>4</sub> and sketch the molecular structure. Indicate whether the molecule is polar or nonpolar, and justify your answer.

Answer:

- (a) all the isotopes have 34 protons but a different number of neutrons in the nucleus.
- (b) 1s<sup>2</sup> 2s<sup>2</sup> 2p<sup>6</sup> 3s<sup>2</sup> 3p<sup>6</sup> 4s<sup>2</sup> 3d<sup>10</sup> 4p<sup>4</sup>  
2 unpaired electrons. 4s [↑↓] 4p [↑↓][↓][↑] Hund's rule indicates that each of the orbitals will be filled with a single electron before it gets paired.
- (c) (i) in Se, the single paired 4p electrons has 1 electron easily removed to create the 3 unpaired 4p orbitals which is energetically favorable; in bromine, the removal of 1 electron still leaves a paired 4p orbital.  
(ii) the shielding effect is stronger in Te and makes it easier to remove an electron (lower ionization energy).

# Bonding and Molecular Structure

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Because F is very electronegative and the molecule is asymmetric with respect to the fluorines, this molecule is polar.

## 2001 D

Account for each of the following observations about pairs of substances. In your answers, use appropriate principles of chemical bonding and/or intermolecular forces. In each part, your answer must include references to both substances.

- Even though NH<sub>3</sub> and CH<sub>4</sub> have similar molecular masses, NH<sub>3</sub> has a much higher normal boiling point (-33°C) than CH<sub>4</sub> (-164°C).
- At 25°C and 1.0 atm, ethane (C<sub>2</sub>H<sub>6</sub>) is a gas and hexane (C<sub>6</sub>H<sub>14</sub>) is a liquid.
- Si melts at a much higher temperature (1,410°C) than Cl<sub>2</sub> (-101°C).
- MgO melts at a much higher temperature (2,852°C) than NaF (993°C).

### Answer:

- NH<sub>3</sub> exhibits hydrogen bonding (H attached to nitrogen, attracted to N in adjacent molecule) between molecules which creates a larger IMF than CH<sub>4</sub> which doesn't exhibit H-bonding, only weak London dispersion forces. More energy is required to overcome this higher IMF in NH<sub>3</sub> and, therefore, has a higher boiling point.
- Both ethane's and hexane's IMF consist mainly of weak London dispersion forces. The greater number of electrons in hexane (50 vs. ethane's 18) creates a greater IMF, enough to make it a liquid at 25°C but for ethane, the fewer electrons make a smaller IMF and that is not strong enough to cause ethane to condense.
- Si forms strong network covalent bonds (4 per atom) to create a high melting solid. The non-polar molecules of Cl<sub>2</sub> (covalent bond, Cl–Cl) do not form strong IMF, only weak London dispersion forces and that makes it easy to melt at a low temperature.
- Magnesium oxide is a Mg<sup>2+</sup>O<sup>2-</sup> ionic compound while sodium fluoride is a Na<sup>+</sup>F<sup>-</sup> ionic compound. The larger ionic charge creates a stronger Coulombic attraction between the anion and cation in MgO and a higher temperature is required to overcome it and melt it.

## 2002 D Required

Use the principles of atomic structure and/or chemical bonding to explain each of the following. In each part, your answer must include references to both substances.

- The atomic radius of Li is larger than that of Be.
- The second ionization energy of K is greater than the second ionization energy of Ca.
- The carbon-to-carbon bond energy in C<sub>2</sub>H<sub>4</sub> is greater than it is in C<sub>2</sub>H<sub>6</sub>.
- The boiling point of Cl<sub>2</sub> is lower than the boiling point of Br<sub>2</sub>.

### Answer:

- Be has 1 more electron in the 2s orbital than Li, not in another larger orbital. Be also has 1 more proton to "pull in" the 2s orbital, making it smaller than Li.

# Bonding and Molecular Structure

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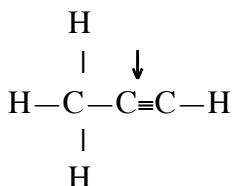
- (b) the second ionization energy ( $\text{IE}_2$ ) removes the second 4s orbital in calcium, leaving a noble gas kernel. The  $\text{IE}_2$  in potassium is removing a very stable 3p orbital electron in its noble gas kernel, which requires a great deal more energy.
- (c) the carbon-to-carbon bond in  $\text{C}_2\text{H}_4$  is a double bond, which is stronger than the carbon-to-carbon single bond in  $\text{C}_2\text{H}_6$ .
- (d)  $\text{Cl}_2$  has 34 electrons and  $\text{Br}_2$  has 70 electrons. more electrons mean greater vander Waals attractions in  $\text{Br}_2$ , more energy to overcome them when it boils and, therefore a higher boiling point than  $\text{Cl}_2$ .

## 2003 D (repeated in organic)

Compound Name	Compound Formula	$\Delta H^\circ_{\text{vap}}$ (kJ mol <sup>-1</sup> )
Propane	$\text{CH}_3\text{CH}_2\text{CH}_3$	19.0
Propanone	$\text{CH}_3\text{COCH}_3$	32.0
1-propanol	$\text{CH}_3\text{CH}_2\text{CH}_2\text{OH}$	47.3

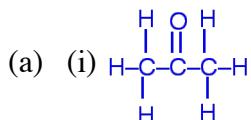
Using the information in the table above, answer the following questions about organic compounds.

- (a) For propanone,
  - (i) draw the complete structural formula (showing all atoms and bonds);
  - (ii) predict the approximate carbon-to-carbon-to-carbon bond angle.
- (b) For each pair of compounds below, explain why they do not have the same value for their standard heat of vaporization,  $\Delta H^\circ_{\text{vap}}$ . (You must include specific information about both compounds in each pair.)
  - (i) Propane and propanone
  - (ii) Propanone and 1-propanol
- (c) Draw the complete structural formula for an isomer of the molecule you drew in, part (a) (i).
- (d) Given the structural formula for propyne below,



- (i) indicate the hybridization of the carbon atom indicated by the arrow in the structure above;
- (ii) indicate the total number of sigma ( $\sigma$ ) bands and the total number of pi ( $\pi$ ) bonds in the molecule

Answer:



(ii)  $120^\circ$

- (b) (i) propane, 26 electrons, molar mass = 44

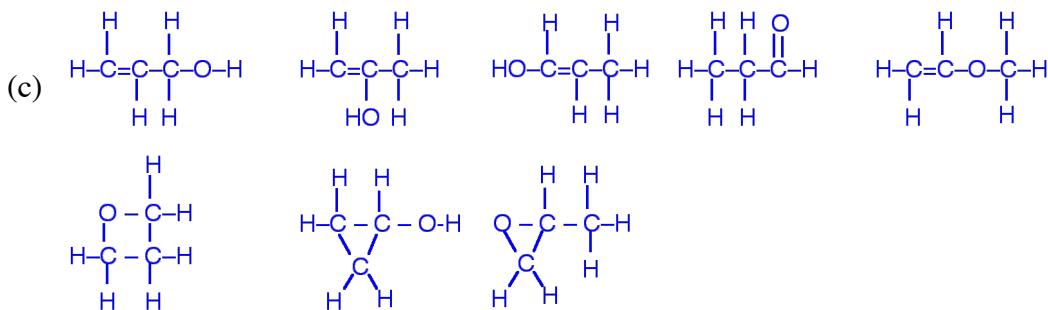
propanone, 32 electrons, molar mass = 58

higher # electrons means larger van der Waal forces, larger molar mass means a slower molecule, the oxygen creates a polar molecule and dipol-dipole interactions

# Bonding and Molecular Structure

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- (ii) 1-propanol has an  $-OH$  which creates a site for hydrogen bonding with other  $-OH$  on adjacent molecules increasing intermolecular forces that must be overcome in order to vaporize the liquid.



- (d) (i)  $sp$   
 (ii) 6 sigma, 2 pi

## 2004 D

Use appropriate chemical principles to account for each of the following observations. In each part, your response must include specific information about both substances.

- At  $25^\circ \text{C}$  and 1 atm,  $\text{F}_2$  is a gas whereas  $\text{I}_2$  is a solid.
- The melting point of  $\text{NaF}$  is  $993^\circ \text{C}$ , whereas the melting point of  $\text{CsCl}$  is  $645^\circ$ .
- The shape of  $\text{ICl}_4^-$  ion is square planar, whereas the shape of  $\text{BF}_4^-$  ion is tetrahedral.
- Ammonia,  $\text{NH}_3$ , is very soluble in water, whereas phosphine,  $\text{PH}_3$ , is only moderately soluble in water.

Answer:

- $\text{F}_2$  is a smaller & lighter molecule than  $\text{I}_2$ , at the same temperature  $\text{F}_2$ , on average is faster than  $\text{I}_2$ . The  $\text{I}_2$  molecule has 106 electrons to the 18 of the  $\text{F}_2$  and, therefore, exhibits a greater vander Waal attraction.
- each ion in  $\text{NaF}$  has a smaller size than the corresponding ion in  $\text{CsCl}$ . This smaller size creates a larger charge density and greater ion Coulombic attraction in the  $\text{NaF}$ , making it harder to melt.
- The  $\text{ICl}_4^-$  ion contains the  $sp^3d^2$  hybridization due to the expanded octet around the central iodine. The chlorides are equatorially bonded in a square around the iodine with the extra pairs of electrons along the axis. The  $\text{BF}_4^-$  ion has  $sp^3$  hybridization which characteristically has the tetrahedral shape.
- ammonia is a more polar molecule than phosphine and can make hydrogen bonds with the solvent, water. This creates a greater solute-solvent attraction and greater solubility.

## 2005 D Required

- Answer the following questions that relate to chemical bonding
- In the boxes provided, draw the complete Lewis structure (electron-dot diagram) for each of the three molecules represented below.

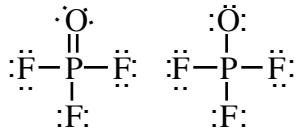
$\text{CF}_4$	$\text{PF}_5$	$\text{SF}_4$

- On the basis of the Lewis structures drawn above, answer the following questions about the particular molecule indicated.

# Bonding and Molecular Structure

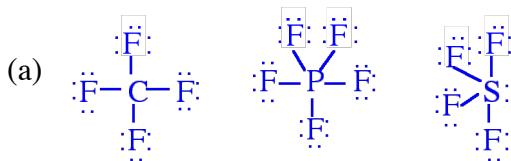
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- (i) What is the F-C-F bond angle in  $\text{CF}_4$ ?
- (ii) What is the hybridization of the valence orbitals of P in  $\text{PF}_5$ ?
- (iii) What is the geometric shape formed by the atoms in  $\text{SF}_4$ ?
- (c) Two Lewis structures can be drawn for the  $\text{OPF}_3$  molecule, as shown below.



- (i) How many sigma bonds and how many pi bonds are in structure 1?
- (ii) Which one of the two structures best represents a molecule of  $\text{OPF}_3$ ? Justify your answer in terms of formal charge.

Answer:



- (b)
- (i)  $109.5^\circ$
  - (ii)  $sp^3d$
  - (iii) see-saw
- (c)
- (i) 4 sigma, 1 pi
  - (ii) structure 1;

In structure 1, oxygen has a formal charge of 0 (6 valence electrons – 6 assigned electrons), each fluorine is 0 (7 valence electrons – 7 assigned electrons), phosphorus is 0 (5 valence electrons – 5 assigned electrons),

In structure 2, oxygen has a formal charge of  $-1$  (6 valence electrons – 7 assigned electrons), each fluorine is 0 (7 valence electrons – 7 assigned electrons), phosphorus is  $+1$  (5 valence electrons – 4 assigned electrons)

According to the electroneutrality rule, the better Lewis structure is the one with the smallest separation of formal charge, *i.e.*, structure 1.

## 2005 D

Use principles of atomic structure, bonding and/or intermolecular forces to respond to each of the following. Your responses must include specific information about all substances referred to in each question.

- (a) At a pressure of 1 atm, the boiling point of  $\text{NH}_3(l)$  is 240 K, whereas the boiling point of  $\text{NF}_3(l)$  is 144 K.
  - (i) Identify the intermolecular forces(s) in each substance.
  - (ii) Account for the difference in the boiling points of the substances.
- (b) The melting point of  $\text{KCl}(s)$  is  $776^\circ\text{C}$ , whereas the melting point of  $\text{NaCl}(s)$  is  $801^\circ\text{C}$ .
  - (i) Identify the type of bonding in each substance.
  - (ii) Account for the difference in the melting points of the substances.
- (c) As shown in the table below, the first ionization energies of Si, P, and Cl show a trend.

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Element	First Ionization Energy (kJ mol <sup>-1</sup> )
Si	786
P	1012
Cl	1251

- (i) For each of the three elements, identify the quantum level (e.g.,  $n = 1$ ,  $n = 2$ , etc.) of the valence electrons in the atom.
- (ii) Explain the reasons for the trend in the first ionization energy.
- (d) A certain element has two stable isotopes. The mass of one of the isotopes is 62.93 amu and the mass of the other isotope is 64.93 amu.
- (i) Identify the element. Justify your answer.
- (ii) Which isotope is more abundant? Justify your answer.

Answer:

(a) (i)

force	NH <sub>3</sub>	NF <sub>3</sub>
London dispersion	+	+
polar attraction	+	+
hydrogen bonding	+	-
ionic attraction	-	-

- (ii) the ability of ammonia to create intermolecular hydrogen bonds, leads to higher amount of energy to separate the molecules by boiling them.
- (b) (i) both compounds have ionic bonding
- (ii) the sodium ion in NaCl is a smaller size than the corresponding potassium ion in KCl. This smaller size creates a larger charge density and greater ion Coulombic attraction in the NaCl, making it harder to melt.
- (c) (i) Si, n = 3; P, n = 3; Cl, n = 3
- (ii) in terms of atomic radius, Si > P > Cl and nuclear charge Cl > P > Si, the smaller and higher charged chlorine atom has the greatest attraction for its electrons than the other two. This means that it takes more energy to remove an electron from chlorine than the other two. The opposite is true for silicon and it should have the smallest value.
- (d) (i) copper; since the atomic mass of an element is the weighted average of its natural isotopes, then the atomic mass of the element must be between 62.93 and 64.94.
- (ii) 62.93; the isotope that is closer to the atom mass of the element is more abundant of the two ( $63.546 - 62.93 = 0.616$ ;  $64.94 - 63.546 = 1.384$ )

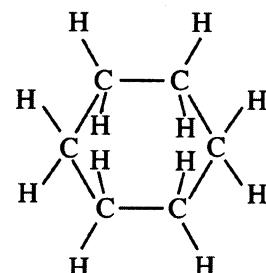
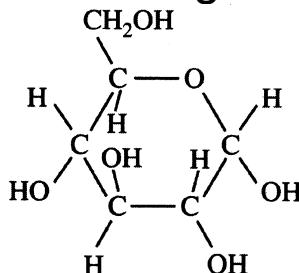
## 2006 D Required

Answer each of the following in terms of principles of molecular behavior and chemical concepts.

- (a) The structures for glucose, C<sub>6</sub>H<sub>12</sub>O<sub>6</sub>, and cyclohexane, C<sub>6</sub>H<sub>12</sub>, are shown below.

# Bonding and Molecular Structure

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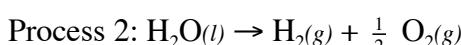


Identify the type(s) of intermolecular attractive forces in

- (i) pure glucose
  - (ii) pure cyclohexane
- (b) Glucose is soluble in water but cyclohexane is not soluble in water. Explain.
- (c) Consider the two processes represented below.



$$\Delta H^\circ = +44.0 \text{ kJ mol}^{-1}$$



$$\Delta H^\circ = +286 \text{ kJ mol}^{-1}$$

- (i) For each of the two processes, identify the type(s) of intermolecular or intramolecular attractive forces that must be overcome for the process to occur.
- (ii) Indicate whether you agree or disagree with the statement in the box below. Support your answer with a short explanation.

When water boils,  $\text{H}_2\text{O}$  molecules break apart to form hydrogen molecules and oxygen molecules.

- (d) Consider the four reaction-energy profile diagrams shown below.

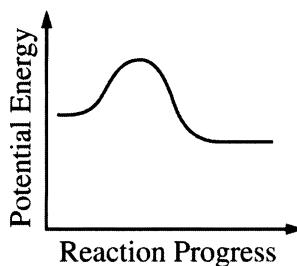


Diagram 1

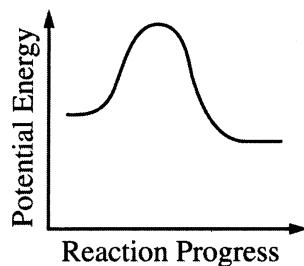


Diagram 2

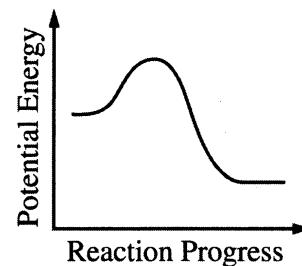


Diagram 3

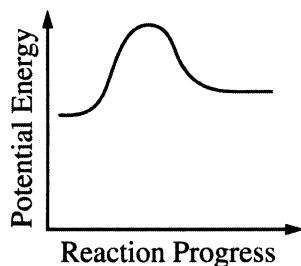


Diagram 4

- (i) Identify the two diagrams that could represent a catalyzed and an uncatalyzed reaction pathway for the same reaction. Indicate which of the two diagrams represents the catalyzed reaction pathway for the reaction.
- (ii) Indicate whether you agree or disagree with the statement in the box below. Support your answer with a short explanation.

Adding a catalyst to a reaction mixture adds energy that causes the reaction to proceed more quickly.

**Answer:**

(a)

London dispersion forces	polar attractions	hydrogen bonding
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# Bonding and Molecular Structure

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(i) pure glucose	+	+	+
(ii) pure cyclohexane	+	-	-

- (b) The hydroxyl groups ( $-\text{OH}$ ) in glucose create polar regions on the molecule, these polar regions can be attracted to the polar water molecules, allowing it to dissolve. Cyclohexane has not such structures and is non-polar and non-water soluble. Like dissolves like.

(c) (i)

	Intermolecular forces			Intramolecular forces
	London dispersion forces	polar attractions	hydrogen bonding	H—O covalent bond
process 1	+	+	+	-
process 2	+	+	+	+

(ii) disagree; there is not enough energy in boiling water (373 K) to break a H—O covalent bond.

(d) (i) diagrams 1 & 2; diagram 1 represents the catalyzed reaction pathway

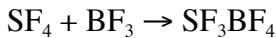
(ii) disagree; a catalyst does not increase the temperature and, therefore, does not increase the amount of energy present in the mixture. It only provides a lower energy pathway (i.e., smaller activation energy requirement) for the reaction to occur.

Answer EITHER Question 7 below OR Question 8. Only one of these two questions will be graded. If you start both questions, be sure to cross out the question you do not want graded. The Section II score weighting for the question you choose is 15 percent.

## 2006 D

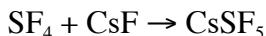
Answer the following questions about the structures of ions that contain only sulfur and fluorine.

(a) The compounds  $\text{SF}_4$  and  $\text{BF}_3$  react to form an ionic compound according to the following equation.



- (i) Draw a complete Lewis structure for the  $\text{SF}_3^+$  cation in  $\text{SF}_3\text{BF}_4$ .
- (ii) Identify the type of hybridization exhibited by sulfur in the  $\text{SF}_3^+$  cation.
- (iii) Identify the geometry of the  $\text{SF}_3^+$  cation that is consistent with the Lewis structure drawn in part (a)(i).
- (iv) Predict whether the F—S—F bond angle in the  $\text{SF}_3^+$  cation is larger than, equal to, or smaller than  $109.50^\circ$ . Justify your answer.

(b) The compounds  $\text{SF}_4$  and  $\text{CsF}$  react to form an ionic compound according to the following equation.

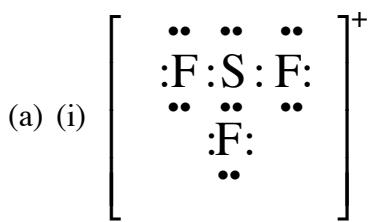


- (i) Draw a complete Lewis structure for the  $\text{SF}_5^-$  anion in  $\text{CsSF}_5$ .
- (ii) Identify the type of hybridization exhibited by sulfur in the  $\text{SF}_5^-$  anion.
- (iii) Identify the geometry of the  $\text{SF}_5^-$  anion that is consistent with the Lewis structure drawn in part (b)(i).
- (iv) Identify the oxidation number of sulfur in the compound  $\text{CsSF}_5$ .

Answer:

# Bonding and Molecular Structure

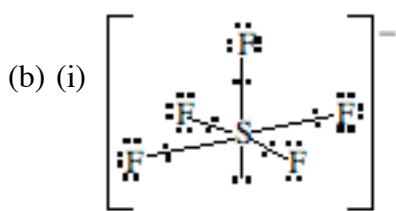
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(ii)  $\text{sp}^3$ .

(iii) pyramidal

(iv) smaller than  $109.50^\circ$ . The lone pair of unbonded electrons occupies more space than the bonded pairs and, therefore, pushed the bonded pairs away and hence, a smaller bond angle than a perfect tetrahedron.



(ii)  $\text{sp}^3\text{d}^2$

(iii) square pyramidal

(iv) S = +4