This page intentionally left blank. Go to next page.
9. 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°C) is a liquid, whereas HCl (normal boiling point = -114°C) is a gas.

(b) Molecules of AsF₃ are polar, whereas molecules of AsF₅ are nonpolar.

(c) The N=O bonds in the NO₂⁻ ion are equal in length, whereas they are unequal in HNO₂.

(d) For sulfur, the fluorides SF₃, SF₄, and SF₆ are known to exist, whereas for oxygen only OF₂ is known to exist.

Scoring Guide

(a) Hydrogen bonding (or dipole-dipole attraction) in HF is greater than it is in HCl

1 point

(Note: only 1 point earned if simply stated that HF has greater intermolecular forces than HCl)

(b) AsF₃ has a trigonal pyramid shape and bond dipoles do NOT cancel (or, asymmetric molecule)

1 point

AsF₅ has a trigonal bipyramid shape and bond dipoles cancel (or, symmetric molecule)

1 point

(Notes: explanation must refer to shape in order to earn point; 1 point earned if only correct Lewis structures are given)

(c) NO₂⁻ has resonance structures

\[ \text{HNO}_2 \text{ has no resonance structures} \]

1 point

\[ \begin{array}{c}
\text{:O=N=O} \\
\text{OR,} \\
\text{one N-O single bond, one N=O double bond}
\end{array} \]

(Notes: 1 point earned if only correct Lewis structures, including resonance for NO₂⁻, are given)

(d) Sulfur uses d orbitals (or expanded octet), oxygen has no d orbitals in its valence shell

1 point

(Sulfur is a larger atom, can accommodate more bonds

2 points

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Question 8

8 points

(a) (i) 2 pts

\[
\begin{align*}
\text{O} & \text{O} \\
\text{O} & \text{O} \\
\text{O} & \text{O} \\
\text{O} & \text{O} \\
\end{align*}
\]

- One point earned for each Lewis electron-dot structure
- Indication of lone pairs of electrons are required on each structure
- Resonance forms of \( \text{CO}_3^{2-} \) are not required

(ii) In \( \text{CO}_2 \), the \( \text{C} \text{-O} \) interactions are double bonds, OR, in \( \text{CO}_3^{2-} \) the \( \text{C} \text{-O} \) interactions are resonance forms (or figures below.) 1 pt

\[
\begin{align*}
\text{O} & \text{O} \\
\text{O} & \text{O} \\
\text{O} & \text{O} \\
\text{O} & \text{O} \\
\end{align*}
\]

The carbon-oxygen bond length is greater in the resonance forms than in the double bonds. 1 pt

- 1st point earned for indicating double bonds are present in \( \text{CO}_2 \) OR resonance occurs in \( \text{CO}_3^{2-} \)
- 2nd point earned for BOTH of the above AND indicating the relative lengths of the bond types

(b) (i) 2 pts

\[
\begin{align*}
\text{F} & \text{F} \\
\text{F} & \text{F} \\
\text{F} & \text{F} \\
\text{F} & \text{F} \\
\end{align*}
\]

- One point earned for each Lewis electron-dot structure
- Lone pairs of electrons are required on each structure
(ii) CF₄ has a tetrahedral geometry, so the bond dipoles cancel, leading to a nonpolar molecule.  

With five pairs of electrons around the central S atom, SF₄ exhibits a trigonal bipyramidal electronic geometry, with the lone pair of electrons. In this configuration, the bond dipoles do not cancel, and the molecule is polar.

![Diagram of CF₄ and SF₄ molecules]

- One point earned for each molecule for proper geometry and explanation
Using principles of chemical bonding and molecular geometry, explain each of the following observations. Lewis electron-dot diagrams and sketches of molecules may be helpful as part of your explanations. For each observation, your answer must include references to both substances.

(a) The bonds in nitrite ion, $\text{NO}_2^-$, are shorter than the bonds in nitrate ion, $\text{NO}_3^-$.  

![Lewis electron-dot diagram for NO₂⁻ and NO₃⁻](image)

According to the Lewis electron-dot diagram, two resonance structures are required to represent the bonding in the $\text{NO}_2^-$ ion. The effective number of bonds between N and O is 1.5.

Three resonance structures are required to represent the bonding in the $\text{NO}_3^-$ ion. The effective number of bonds between N and O is 1.33.

The greater the effective number of bonds, the shorter the N–O bond length.

(b) The $\text{CH}_2\text{F}_2$ molecule is polar, whereas the $\text{CF}_4$ molecule is not.

The molecular geometry in both $\text{CH}_2\text{F}_2$ and $\text{CF}_4$ is tetrahedral (or the same). The C-F bond is polar. In $\text{CF}_4$, the molecular geometry arranges the C-F dipoles so that they cancel out and the molecule is nonpolar. The C-H bond is less polar than the C-F bond. The two C-H dipoles do not cancel the two C-F dipoles in $\text{CH}_2\text{F}_2$.  

| 1 point earned for discussing the similarity in molecular geometry |
| 1 point earned for discussing the relationship between molecular geometry and the C-H and C-F bond dipoles |
Question 6 (cont’d.)

(c) The atoms in a $\text{C}_2\text{H}_4$ molecule are located in a single plane, whereas those in a $\text{C}_2\text{H}_6$ molecule are not.

| The carbon atoms in $\text{C}_2\text{H}_4$ have a molecular geometry around each carbon atom that is trigonal planar ($\text{AX}_3$), so all six atoms are in the same plane. The carbon atoms in $\text{C}_2\text{H}_6$ have a molecular geometry that is tetrahedral ($\text{AX}_4$), so the atoms are not all in the same plane. | 1 point earned for the bonding of the carbon atoms  
1 point earned for the structure |
| OR | The carbon-carbon double bond in $\text{C}_2\text{H}_4$ results in a planar molecule whereas the carbon-carbon single bond in $\text{C}_2\text{H}_6$ results in a non-planar (tetrahedral) site at each carbon atom. |

(d) The shape of a $\text{PF}_5$ molecule differs from that of an $\text{IF}_5$ molecule.

| In $\text{PF}_5$, the molecular geometry is trigonal bipyramidal because the phosphorus atom has five bonding pairs of electrons and no lone pairs of electrons. $\text{IF}_5$ has square pyramidal molecular geometry. The central iodine atom has five bonding pairs of electrons and one lone pair of electrons. The presence of the additional lone pair of electrons on the central iodine atom means the molecular geometry is different. | 1 point earned for discussing the difference made by the lone pair of electrons in $\text{IF}_5$ and how it affects the geometry of the two molecules |

(e) $\text{HClO}_3$ is a stronger acid than $\text{HClO}$.

| According to the formula for $\text{HOCl}$ and $\text{HOClO}_2$, there are two additional terminal, electronegative oxygen atoms attached to the central chlorine atom. These additional terminal oxygen atom stabilize the negative charge on the anion $\text{ClO}_3^-$ compared to $\text{ClO}^-$. The result is to reduce the electrostatic attraction between the $\text{H}^+$ and $\text{ClO}_x^-$. OR | 1 point earned for discussing the importance of the electronegativity of the terminal oxygen atoms in the two structures and/or the enhanced stability of the chlorate vs. the hypochlorite ion |
| The two additional terminal electronegative O atoms bonded to the chlorine atom of $\text{ClO}_3^-$ pull electron density away from the central chlorine atom. The net result is to weaken the H-O bond. Since $\text{HOCl}$ has no additional terminal O atoms, its H-O bond is stronger. The weaker the H-O bond, the stronger the acid. |
Question 8

<table>
<thead>
<tr>
<th>Compound Name</th>
<th>Compound Formula</th>
<th>$\Delta H_{vap}^o$ (kJ mol$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propane</td>
<td>CH$_3$CH$_2$CH$_3$</td>
<td>19.0</td>
</tr>
<tr>
<td>Propanone</td>
<td>CH$_3$COCH$_3$</td>
<td>32.0</td>
</tr>
<tr>
<td>1-propanol</td>
<td>CH$_3$CH$_2$CH$_2$OH</td>
<td>47.3</td>
</tr>
</tbody>
</table>

8. 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);

```
  H   O   H
  | \ | | |
H---C---C---H
  | \ | |
H   H
```

1 point for complete and correct structural formula

(ii) predict the approximate carbon-to-carbon-to-carbon bond angle.

The C – C – C bond angle is 120°

1 point for bond angle

(b) For each pair of compounds below, explain why they do not have the same value for their standard heat of vaporization, $\Delta H_{vap}^o$. (You must include specific information about both compounds in each pair.)

(i) Propane and propanone

The intermolecular attractive forces in propane are dispersion forces only. The IMFs in propanone are dispersion and dipole-dipole. Since the intermolecular attractive forces differ in the two substances, the enthalpy of vaporization will differ.

1 point for correctly identifying the IMFs for each substance
(ii) Propanone and 1-propanol

| The intermolecular attractive forces in 1-propanol are dispersion forces and hydrogen-bonding. The IMFs in propanone are dispersion and dipole-dipole. Since the intermolecular attractive forces differ in the two substances, the enthalpy of vaporization will differ. | 1 point for correctly identifying the IMFs for each substance |

(c) Draw the complete structural formula for an isomer of the molecule you drew in part (a) (i).

| H H O
  |   |
  H—C—C—C—H
  |   |
  H H |

1 point for correct, complete structural formula

(d) Given the structural formula for propyne below,

| H
  |   |
  H—C—C≡C—H
  |   |
  H |

(i) indicate the hybridization of the carbon atom indicated by the arrow in the structure above;

| sp hybridization | 1 point for correct hybridization |

(ii) indicate the total number of sigma (σ) bonds and the total number of pi (π) bonds in the molecule.

| 6 sigma bonds 2 pi bonds | 1 point for correct number of sigma bonds 1 point for correct number of pi bonds |