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McCord CH301

Exam 4

Nov 14, 2017

Remember to refer to the Periodic Table handout that is separate from this exam copy.

NOTE: Please keep this exam copy intact (all pages still stapled including this cover page). You must turn in ALL the materials that were distributed. This means that you turn in your exam copy (name and signature included), bubble sheet, periodic table handout, and all scratch paper. Please also have your UT ID card ready to show as well.

This print-out should have 20 questions. Multiple-choice questions may continue on the next column or page – find all choices before answering.

001 5.0 points

A solid sample dissolves in water but does not conduct electricity. The sample melts at 433K. Which type of solid is this?

- 1. Metallic solid
- 2. Ionic solid
- 3. Covalent network solid
- 4. Molecular solid correct

Explanation:

Molecular solids have relatively low melting points and do not conduct electricity.

002 5.0 points

What types of intermolecular forces are present in a homogenous mixture of the molecule shown below?

- ${f 1.}$ dispersion forces, dipole-dipole ${f correct}$
- 2. dispersion forces only
- 3. hydrogen bonding only
- **4.** dispersion forces, dipole-dipole, hydrogen bonding
 - 5. dispersion forces, hydrogen bonding
- **6.** dispersion forces, dipole-dipole, ion-dipole

Explanation:

The molecule shown has polar bonds. Therefore, it has dipole-dipole interactions. All molecules are capable of dispersion forces.

003 5.0 points

Rank the following substances in order of freezing point: Ne, Rn, Kr, He

- 1. Rn < He < Ne < Kr
- **2.** Kr < Ne < Rn < He
- 3. He < Ne < Kr < Rn correct
- 4. Kr < Ne < He < Rn
- 5. He < Ne < Rn < Kr

Explanation:

When determining the relative IMFs within a group, you should rank based on polarizability (molecular weight). Therefore, the ranking is: He < Ne < Kr < Rn

004 5.0 points

Arrange the following substances in order from highest to lowest melting point:

 SiO_2 , CH_3CH_2OH , RbF, $CaSO_4$

- 1. $SiO_2 > CaSO_4 > RbF > CH_3CH_2OH$ correct
 - 2. $CH_3CH_2OH > CaSO_4 > RbF > SiO_2$
 - 3. $SiO_2 > RbF > CaSO_4 > CH_3CH_2OH$
 - 4. $SiO_2 > CaSO_4 > CH_3CH_2OH > RbF$
 - 5. $CH_3CH_2OH > RbF > CaSO_4 > SiO_2$

Explanation:

SiO₂ is a network covalent solid. Both CaSO₄ and RbF are ionic solids, but CaSO₄ has a greater charge density. CH₃CH₂OH is a molecular solid. Thus, arranging the compounds in order from highest to lowest melting point, we get:

$$\mathrm{SiO}_2 > \mathrm{CaSO}_4 > \mathrm{RbF} > \mathrm{CH}_3\mathrm{CH}_2\mathrm{OH}$$
 .

005 5.0 points

Which of the following best describes why H_2 Te has a higher boiling point than H_2 S?

- 1. H_2 Te is more polar than H_2S
- **2.** H_2S is larger than H_2Te
- **3.** H₂Te is capable of hydrogen bonding
- 4. H_2S is more polarizable than H_2Te
- 5. H_2 Te is more polarizable than H_2 S correct
 - **6.** None of the above

Explanation:

Both H_2 Te and H_2 S have similar polar molecular structures. However, H_2 Te will be more polarizable because Te is larger than S, resulting in a higher BP.

006 5.0 points

State the dominant intermolecular force in each compound, respectively:

$$CBr_4$$
, NH_3 , CH_3F , $C_{12}H_{14}$

- 1. dipole-dipole, hydrogen-bonding, dipole-dipole, hydrogen-bonding
- 2. dipole-dipole, hydrogen-bonding, hydrogen-bonding, dispersion
- **3.** dispersion, hydrogen-bonding, dipole-dipole, dispersion **correct**
- **4.** dipole-dipole, dispersion, hydrogenbonding, dispersion
- **5.** dispersion, hydrogen-bonding, hydrogen-bonding, dispersion
- **6.** dispersion, dipole-dipole, hydrogenbonding, dispersion

Explanation:

CBr₄ consists of polar bonds but the molecule is overall non-polar. Therefore, CBr₄ is dominated by dispersion forces. NH₃ is capable of hydrogen bonding. CH₃F looks like it could have hydrogen bonding, but the fluorine is not connected to a hydrogen. Therefore, CH₃F is dominated by dipole-

dipole interactions. $C_{12}H_{14}$ is an organic, non-polar molecule dominated by dispersion forces.

007 5.0 points

In a molecular orbital diagram for C_2 , the LUMO refers to the

- 1. σ_{2s}^*
- **2.** π_{2p}^*
- **3.** σ_{2p}^*
- 4. σ_{2s}
- 5. σ_{2p} correct
- **6.** π_{2p}

Explanation:

 C_2 has eight valence electrons.

008 5.0 points

Which of the following statements is true regarding molecular orbital theory?

- 1. Energy is conserved when pure atomic orbitals combine to form bonding and anti-bonding orbitals **correct**
- 2. Antibonding orbitals lend energetic stability to the overall molecule
- **3.** All diatomic molecules with an even number of total electrons are diamagnetic
- **4.** Molecular orbital theory can only depict the bonding of homonuclear diatomic molecules

Explanation:

Not all diatomic molecules with an even number of total electrons are diamagnetic. An example of this is O_2 . Molecular orbital theory is advantageous because complex diagrams can depict the bonding of heteronuclear molecules of all sizes. The only true statement is: Energy is conserved when pure atomic orbitals combine to form bonding and antibonding orbitals

009 5.0 points

A liquid poured into a graduated cylinder forms a concave meniscus. Which of the following statements is true regarding this liquid sample?

- 1. The meniscus is formed to minimize the attractive forces between the liquid molecules and the walls of the container
- 2. The meniscus is concave because the adhesive forces are lower in energy than the cohesive forces **correct**
- **3.** The meniscus forms to maximize the potential energy of the liquid
- 4. The liquid molecules on the surface are more attracted to each other than the walls of the container

Explanation:

A meniscus is always formed to maximize attractions and minimize the potential energy. In this case, a concave meniscus is formed by adhesive forces between liquid molecules and the walls of the container.

010 5.0 points

Consider the MO diagram for F_2^+ . What is the bond order? Is F_2^+ paramagnetic or diamagnetic?

- 1. 2, paramagnetic
- 2. 1, diamagnetic
- **3.** 2.5, paramagnetic
- 4. 2, diamagnetic
- 5. 2.5, diamagnetic
- 6. 1.5, diamagnetic
- 7. 1.5, paramagnetic correct

Explanation:

 F_2^+ has 8 bonding electrons and 5 antibonding electrons.

BO =
$$\frac{1}{2}$$
[(bonding e⁻) - (antibonding e⁻)]
= $\frac{1}{2}$ (8 - 5) = 1.5

 F_2^+ has unpaired electrons and is therefore paramagnetic.

011 5.0 points

Which of the following has the lowest surface tension?

- 1. OH
- 2. HO OH
- 4. NH₂
- 5. correct

Explanation:

The molecule that has the lowest surface tension will have dispersion forces only. Of the molecules shown, the molecule will the weakest IMF's will exhibit the most hydrogen bonding.

Comparing organic molecules, more double bonds correspond to a smaller HOMO-LUMO gap. For example, β -carotene has a pi bond network of 11 bonds and absorbs light in the visible spectrum. An advantage of this is that organic molecules can be selectively synthesized to absorb at specific wavelengths. Based on this information, if you wanted to synthesize a sunscreen that absorbs light in the UV range you would want

- 1. to absorb at longer wavelengths than β -carotene
- **2.** a larger network of pi bonds than β -carotene
 - 3. none of the above
- **4.** a smaller network of pi bonds than β -carotene **correct**
- **5.** a smaller HOMO-LUMO gap than β -carotene

Explanation:

The question stem states that more double bonds (a larger network of pi bonds) corresponds to a smaller HOMO-LUMO gap. To absorb at the UV range, you would need a large HOMO-LUMO gap. This means you will need fewer double bonds or a small network of pi bonds.

013 5.0 points

The instability of a bond associated with antibonding orbitals is best explained by...

- 1. Molecular Orbital Theory correct
- 2. Valence Bond Theory

Explanation:

Molecular orbital theory gives us insight into bonding and anti-bonding orbitals.

014 5.0 points

A theoretical central atom has three bonding regions and two lone pair(s). What is the hybridization of this atom?

- $1. \mathrm{sp}^3$
- 2. $\mathrm{sp}^3\mathrm{d}$ correct
- **3.** sp
- **4.** $sp^{3}d^{2}$
- **5.** sp^2

Explanation:

To determine the valence bond hybridization of a molecule, you must consider all regions of electronic density. Five total electronic regions is sp³d.

015 5.0 points

The following structures are constitutional isomers, meaning they share the same molecular formula but the atoms are connected differently.

n-heptane:

3-ethyl pentane:



Which of the following best explains why n-heptane has a higher boiling point than 3-ethyl pentane?

- 1. 3-ethyl pentane is an overall larger molecule, which interferes with intermolecular forces
- **2.** *n*-heptane has a greater surface area of overlap that can polarize to form instantaneous dipoles **correct**
- **3.** *n*-heptane is an overall larger molecule, which interferes with intermolecular forces

- 4. Branched molecules allow dispersion forces to be more tightly packed than linear molecules
 - **5.** *n*-heptane is a more polar molecule

Explanation:

Dispersion forces have a very short range of interaction. Therefore, a molecule that can be more tightly packed will have a higher boiling point. Linear molecules, such as *n*-heptane have more regions of electron density that can form instantaneous dipoles (dispersion forces).

016 5.0 points

Consider the valence bond configuration for a central atom:

$$\begin{array}{c|c} \uparrow \downarrow & \uparrow & \uparrow \\ \hline sp^3 & \end{array}$$

This central atom belongs to which of the following molecules?

1.
$$\left[\ddot{\mathbf{O}} = \ddot{\mathbf{N}} - \ddot{\mathbf{O}} : \right]$$

2.
$$\begin{array}{ccc} H - \overset{\cdots}{P} - H \\ & & \text{correct} \\ H \end{array}$$

3.
$$\ddot{O} = C = \ddot{O}$$

Explanation:

The valence bond configuration shows one lone pair and three sigma bonds. This is PH₃.

017 5.0 points

You are able to extract two saturated hydrocarbons, octane and decane, from a liquid organic mixture. Both molecules are linear hydrocarbons (i.e. there is no branching). Compared to octane, decane appears more viscous. You can conclude that decane also has:

- I. a higher molecular weight
- II. a smaller surface area
- III. a higher boiling point
- IV. a higher vapor pressure
- 1. III only
- 2. I, II and III only
- **3.** none of the above
- 4. I and III only correct
- **5.** II and III only
- 6. I, II, III, and IV

Explanation:

Viscosity is a qualitative way of determining the relative strength of IMF's. Because decane is more viscous, we know that it has stronger IMF's. Therefore, it will be expected to have a higher boiling point and a lower vapor pressure. A more branched structure will weaken the overall IMFs, so we cannot make this conclusion with the information provided. Because both structures are saturated hydrocarbons, we can conclude that decane has a higher MW.

018 (part 1 of 2) 5.0 points

Consider the following organic line structure:

According the valence bond theory, how many sigma and pi bonds are in the organic structure above?

- 1. 3 σ and 12 π
- **2.** 14 σ and 3 π
- **3.** 18 σ only
- **4.** 18 σ and 3 π
- 5. 15 σ and 3 π correct

Explanation:

Accounting for the "understood" hydrogen atoms, there are a total of 12 single bonds and 3 double bonds. This is a total of 15 σ and 3 π .

019 (part 2 of 2) 5.0 points

For the structure shown above, what is the hybridization around oxygen?

- 1. sp^2
- 2. None of the above
- 3. sp^3 correct
- **4.** sp
- **5.** sp^3d^2
- **6.** $sp^{3}d$

Explanation:

In this structure, oxygen has four regions of electron density, corresponding to a sp^3 hybridization.

020 5.0 points

Name the bond in carbon dioxide (CO_2) that

overlaps above and below the internuclear axis.

- 1. σ_{sp-sp}
- **2.** σ_{sp^2-2p}
- 3. $\sigma_{sp^2-sp^2}$
- **4.** $\pi_{sp^2-sp^2}$
- **5.** σ_{2p-2p}
- **6.** $\pi_{sp^3-sp^3}$
- 7. π_{2p-2p} correct
- 8. σ_{sp^2-sp}

Explanation:

Pi bonds consist of overlapping p orbitals off the internuclear axis. For carbon dioxide, the 2p atomic orbitals overlap above and below the internuclear axis.