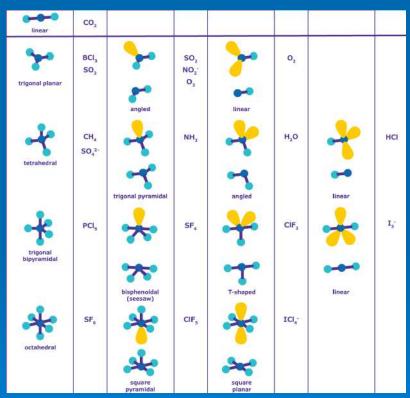
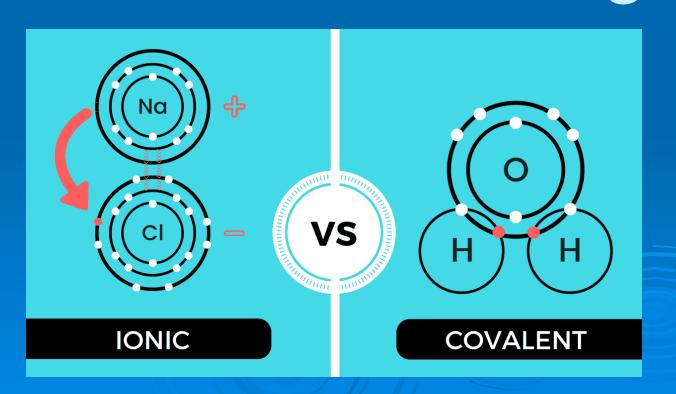
AP Chemistry Chapter 8 and 9 Jeopardy



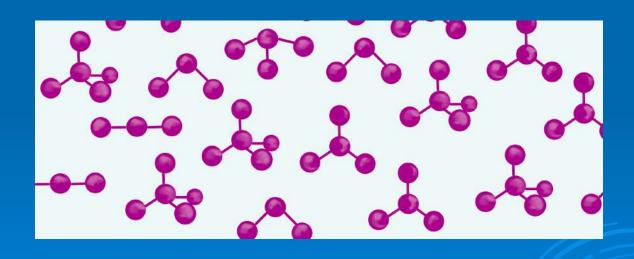
Jennie L. Borders

Round 1 – Chapter 8 Basic Concepts of Chemical Bonding



Lattice Energy	Polarity	Lewis Dot Structures	Formal Charge	Resonance	Surprise
100	100	100	100	100	100
200	200	200	200	200	200
300	300	300	300	300	300
400	400	400	400	400	400
500	500	500	500	500	500

Round 2 – Chapter 9 Molecular Geometry and Bonding Theories



Click to go to Round 2

Define lattice energy.

Lattice energy is the energy required to completely separate a mole of a solid ionic compound into its gaseous ions.

Explain the following trend in lattice energy: BaO > KF.

BaO has a larger magnitude of charge (+2 and -2), KF only has +1 and -1.

Explain the following trend in lattice energy: CaF₂ > BaF₂.

CaF₂ has a smaller cation, so lattice energy is larger.

Arrange the following compounds in order of lowest to highest lattice energy: ScN, KBr, MgS, NaCl.

KBr, NaCl, MgS, ScN

The table below shows the melting points of $MgO_{(s)}$ and $NaF_{(s)}$. Which of the following best helps explain why the melting point of $MgO_{(s)}$ is much higher than that of

Compound	Melting Point (°C)		
MgO	2852		
NaF	993		

NaF_(s)?

- a. The mass of F- ions is greater than that of O2- ions.
- b. The mass of Mg²⁺ ions is greater than that of Na⁺ ions.
- c. The difference between the electronegativity values of Mg and O is smaller than the difference between the values of Na and F.
- d. The charges of Mg²⁺ and O²⁻ ions are greater than those of Na⁺ and F⁻ ions.

Of the following pair, which one is a molecular substance and which one is an ionic substance: PbCl₄ or RbCl.

PbCl₄ = molecular RbCl = ionic

Of the following compounds, which is the most ionic?

a.SiCl₄

b.BrCl

c.PCI₃

d.Cl₂O

e.CaCl₂

9.

Of the following single bonds, which is the LEAST polar?

a.N - H

b.H - F

c.O - F

d.I - F

e.O – H

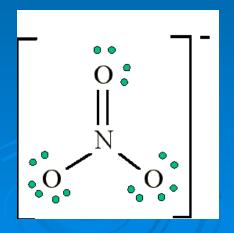
3.

Draw the dipole moment for the following bond O – P.

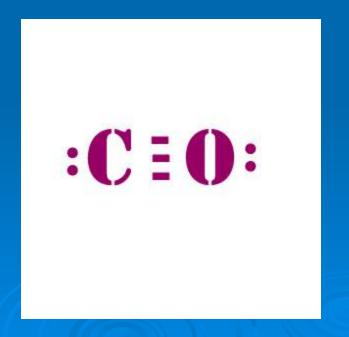


Draw the dipole moments for the following molecule: NO₃-.

3 resonance structures, dipoles point toward the O's



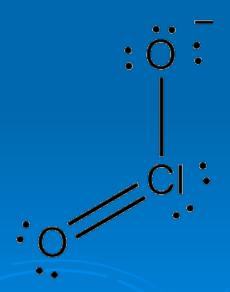
Draw the Lewis dot structures for CO.



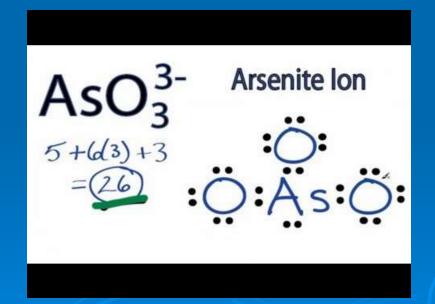
Draw the Lewis dot structure for C₂H₂.

$$H \longrightarrow C \equiv C \longrightarrow H$$

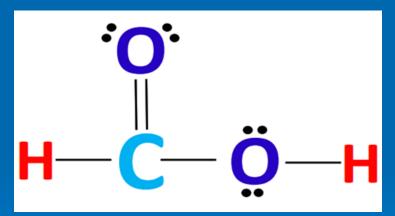
Draw the Lewis dot structure for CIO₂-.



Draw the Lewis dot structure for AsO₃³-.



 a. Complete the Lewis electron-dot diagram for HCOOH. Show all bonding and nonbonding valence electrons.

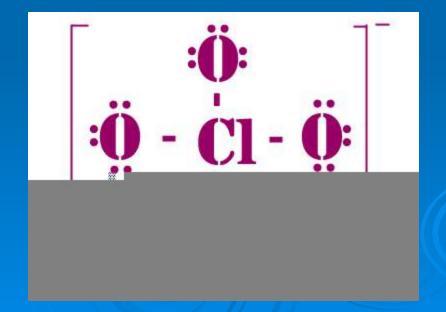


Yes, the oxidation number of C changes from +2 to +4 and H changes from +1 to 0.

b. Is the following a redox reaction? Justify your answer.

$$HCOOH_{(aq)} \rightarrow H_{2(g)} + CO_{2(g)}$$

Determine the formal charges on the atoms of ClO₄-.

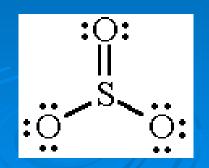


Determine the formal charges on the atoms of NO⁺.

Formal charge on N = 0 and O = +1

Determine the formal charges on the atoms of SO₃. (make sure all atoms follow the octet rule)

Formal charge on S = +2, on the double bonded O = 0, and on the 2 single bonded O = -1



Two possible Lewis electron-dot diagrams for CO₂ are shown below. Explain in terms of formal charges why diagram Z is the better diagram. Also, identify the hybridization of the valence orbitals of the C atom in the CO₂ molecule in diagram Z.

Diagram X

:O-C≡O:

Diagram Z

O=C=O

Diagram Z has a formal charge of zero on every atom, so it is preferred. The hybridization is sp.

Two possible Lewis electron-dot diagrams for fulminic acid, HCNO, are shown below.

$$H-C\equiv N-\ddot{O}$$
: $H-\ddot{C}=N=\ddot{O}$:

Explain why the diagram on the left is the better representation for the bonding in fulminic acid. Justify your choice based on formal charges.

Left: C = 0 and O = -1. Right: C = -1 and O = 0. The diagram on the left is better because it puts the negative formal charge on oxygen, which is more electronegative than carbon.

Draw the resonance structures for NO₂-.

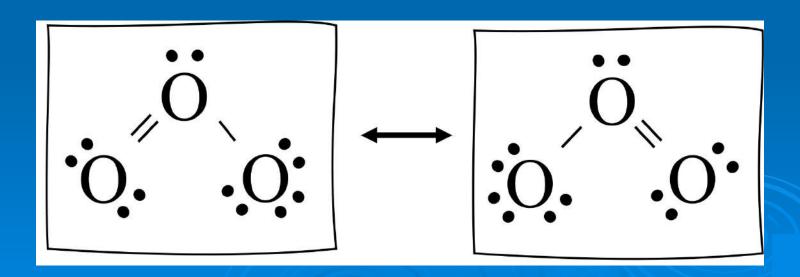


Draw the resonance structures for NO_2^+ .

Example: nitronium ion -
$$NO^+_2$$

 $O=N=O$
 $\longrightarrow :O=N-O:$
 $\longrightarrow :O-N=O:$

The O₃ molecule has a central oxygen atom bonded to two outer oxygen atoms that are not bonded to one another. Draw the Lewis electrondot diagram of the O₃ molecule. Include all valid resonance structures.



Predict the ordering of the C – O bond lengths in CO, CO₂, and CO_3^{2-}



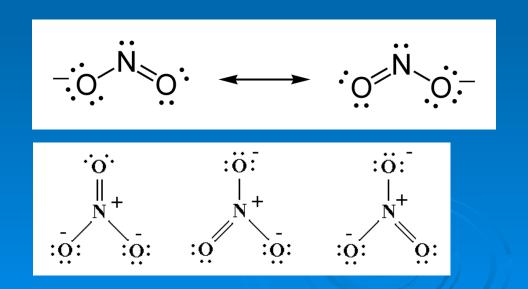
 $: \mathbb{C} = \mathbb{O}:$ CO, CO₂, CO₃² $\ddot{O} = C = \ddot{O}$

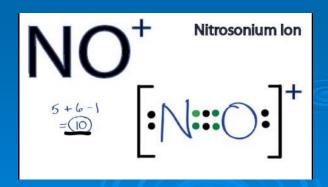
$$\ddot{O} = C = \ddot{O}$$

$$\begin{bmatrix} : \ddot{O} : \\ & \downarrow \\ & \downarrow$$

Based on the Lewis structures, predict the ordering of N – O bond lengths in NO₂-, NO₃-, and NO+.

 NO^+ , NO_2^- , NO_3^-





The table below provides information about two types of steel. Which of the following best helps explain why high-carbon steel is more rigid than low-carbon steel?

Type of Steel	% Carbon	Characteristics	Uses
Low-carbon steel	< 0.2%	Malleable and ductile	Chains and nails
High-carbon steel	0.6 - 1.5%	Hard and brittle	Cutting tools

- a. Elemental carbon is harder than elemental iron.
- b. The additional carbon atoms within the alloy make the high-carbon steel less dense.
- c. The additional carbon atoms within the alloy increase the thermal conductivity of the high-carbon steel.
- d. The additional carbon atoms within the alloy make it more difficult for the iron atoms to slide past one another.

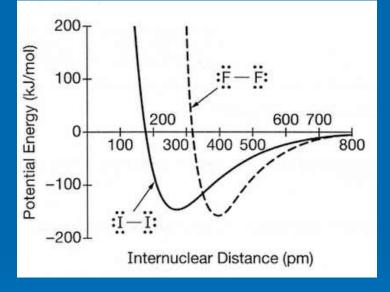
White gold is a common alloy of gold and palladium that is often used in jewelry. The atomic radii of the metals are given in the table below. A particular ring is made from an alloy that is 75 mole percent and 25 mole percent palladium. Draw a particle-level diagram of the solid alloy consisting of 12 atoms with a representative proportion of atom types. Your diagram should clearly indicate whether the alloy is interstitial of substitutional. Use empty circles for gold and shaded circles for palladium.

Element	Atomic Radius (pm)		
Au	135		
Pd	140		

12 circles of similar size. 3 shaded and 9 empty.

A student incorrectly sketched the potential energy curve for two fluorine atoms, shown by the

dotted line.



Explain the error with the student's sketch for

The minimum of the curve for F₂ should be to the left of that for I₂ since the atomic radius of F is smaller than the atomic radius of I, leading to the F-F bond being shorter than the I-I bond.

The molecular formulas of diatomic bromine, chlorine, fluorine, and iodine are written below. Pick the formula of the molecule that has the longest bond length. Justify your answer in terms of atomic structure. Br₂, Cl₂, F₂, l₂

I₂ has the longest bond length because the radius of the I atoms is greater than the radii of the other halogen atoms. Thus, the distance between the nuclei of atoms in I₂ is greater than it is in smaller halogens.

To make Au stronger and harder, it is often alloyed with other metals, such as Cu and Ag. Consider two alloys, one of Ag and Cu and one of Au and Ag, each with the same mole fraction of Au. If the Au/Cu alloy is harder than the Au/Ag alloy, then which of the following is the best explanation based on the information in the table below?

Element	Metallic Radius (pm)	Melting Point (°C)	Common Oxidation State
Au	144	1064	1+, 3+
Cu	128	1085	1+, 2+
Ag	144	961	1+

C.

- a. Cu has two common oxidation states, but Ag has only one.
- b. Cu has a higher melting point than Au has, but Ag has a lower melting point than Au has.
- c. Cu atoms are smaller than Ag atoms, thus they interfere with the displacement of atoms in the alloy.
- d. Cu atoms are less polarizable than are Au or Ag atoms, thus Cu has weaker interparticle forces.

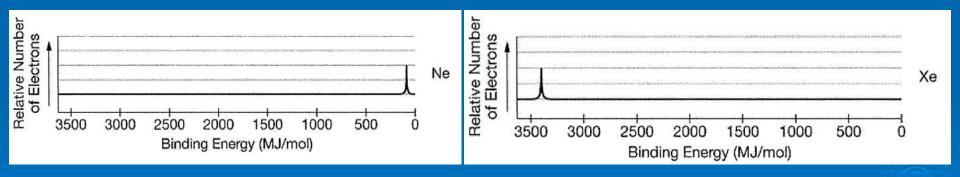
Free Response (Must do all 5)	Bond Angles	VSEPR Model	Polarity	Hybrid Orbitals	Multiple Bonds
200	200	200	200	200	200
400	400	400	400	400	400
600	600	600	600	600	600
800	800	800	800	800	800
1000	1000	1000	1000	1000	1000

Free Response 200

Using principles of atomic structure, explain why the atomic radius of Xe is less than that of Te.

Xe has 54 p⁺ and Te has 52 p⁺. They both have ve⁻ on the 5th energy level. Since Xe has more p⁺, then its ve⁻ have a greater attraction for the nucleus which causes the ve⁻ to be pulled in closer resulting in Xe having a smaller atomic radius.

Photoelectron spectroscopy data for the 1s sublevel of Ne and the 1s sublevel of Xe are represented below. In terms of Coulomb's law and atomic structure, explain why the peak for Ne is positioned so far to the right of the peak for Xe.



Xe has 54 p⁺ and Ne has 10 p⁺. Since Xe has more protons, its 1s e⁻ have more attraction for the nucleus which pulls them closer to the nucleus causing a larger binding energy.

Three compounds that contain an element from Group 18 are XeOF₄, XeO₂F₂, and XeF₄. Possible Lewis electron dot diagrams for the compounds are shown below.

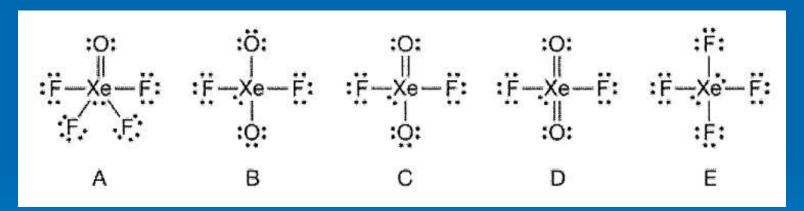
Which structure, B, C, or D, minimizes the formal charges for each atom in XeO₂F₂? Justify your choice.

Structure D because every atom has a formal charge of zero.

A student proposes that the energy of the Xe – to – F bond in the structure in diagram B is equal to that of the Xe – to – F bond in the structure in diagram A. Do you agree or disagree with the student's claim? Justify your answer.

Agree. The Xe – F bond in structure B and A are both since bonds, so they require the same amount of energy to break.

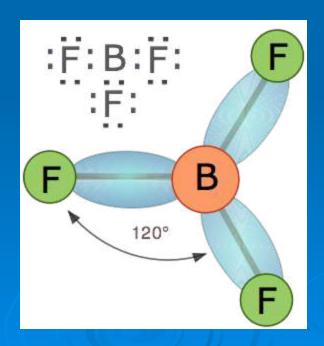
The dipole moment of the XeOF₄ molecule (diagram A) is nonzero. Identify its molecular geometry and explain how its geometry and the polarity of its bonds results in a nonzero dipole moment.



The molecular geometry is square pyramidal. Since there is a lone pair of electrons on Xe, the dipoles cannot cancel out.

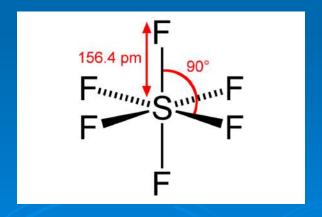
What are the bond angles in the following molecule: BF₃.

Bond angles = 120°



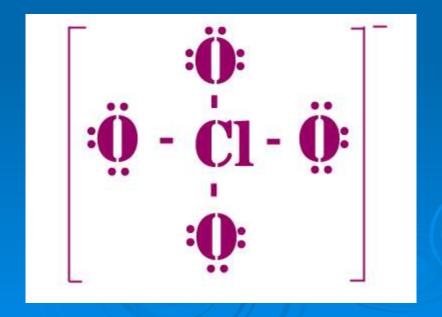
What are the bond angles in the following molecule: SF₆.

Bond angles = 90°



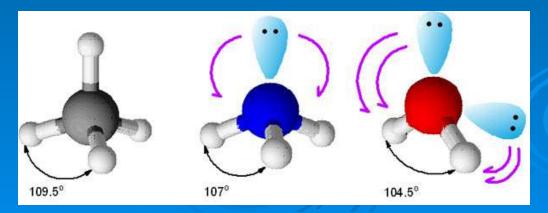
What are the bond angles in the following molecule: CIO₄-.

Bond angles = 109.5°



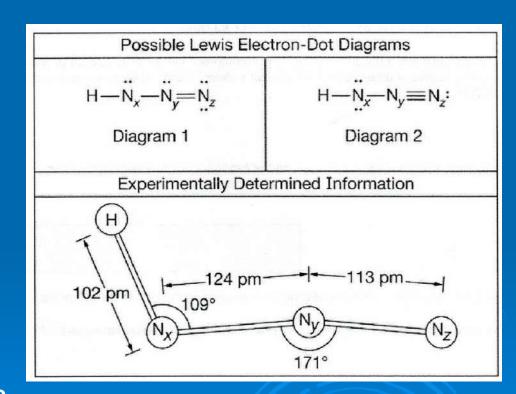
Explain the bond angle differences in CH₄, H₂O, and NH₃.

The bond angles decreases as more lone pairs are added since lone pairs take up more space than bonded electrons.

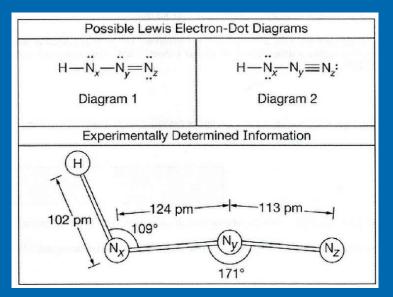


Two possible Lewis electron-dot diagrams and experimentally determined information for HN₃ are shown below.

- a. Explain why diagram 2 is consistent with the experimentally determined bond lengths shown above.
- b. Use VSEPR theory to explain why Diagram 1 is NOT consistent with the experimentally determined bond angles around N_y.



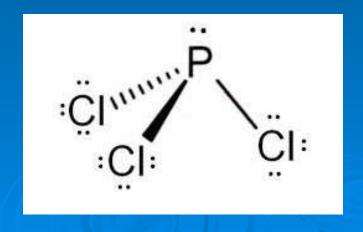
Bond Angles 1000 Answer



- a. Diagram 2 has a single bond between N_x and N_y and a triple bond between N_y and N_z . Triple bonds are shorter than single bonds, so this is consistent with the data.
- b. VSPER shows that 2 bonding and 1 lone pair around N_y make a bent geometry with a bond angle of 120°. This is inconsistent with the 171° bond angle in the data.

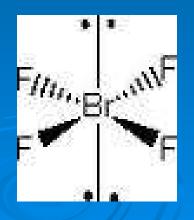
Draw a complete Lewis electron-dot diagram for PCl₃. What is the shape of the PCl₃ molecule?

MG = trigonal pyramidal



What is the molecular geometry and electron domain geometry of the following molecule: BrF₄.

MG = square planar EDG = octahedral



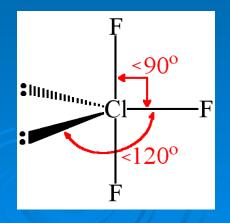
Which of the following has molecules with a pyramidal shape?

- a. NH_{3(g)}
- b. BH_{3(g)}
- c. $H_{2(g)}$
- d. $H_2S_{(g)}$
- e. HBr_(g)

a.

What is the molecular geometry and electron domain geometry of the following molecule: CIF₃.

MG = T-shaped EDG = trigonal bipyramidal

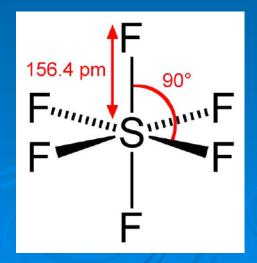


An unknown gas, D, reacts with fluorine gas to form the compound $DF_{4(g)}$. A student claims that if element D is in group 18, then the molecular geometry of $DF_{4(g)}$ is tetrahedral. Do you agree or disagree with the student? Justify your answer in terms of VSEPR theory.

Disagree. If D is in group 18, then DF₄ has a total of 36 ve⁻. This would make the geometry square planar instead of tetrahedral because the 2 extra lone pairs would go on D.

Is the following molecule polar or nonpolar: SF₆. (If polar, show the overall dipole moment.)

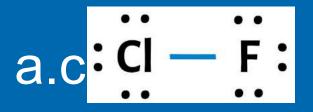
Nonpolar – dipoles all point towards the F's and cancel



Is the following molecule polar or nonpolar: IF. (If polar, show the overall dipole moment.)

Polar – dipole points towards the F

Which of the following Lewis electron-dot diagrams represents the molecule that is the most polar?

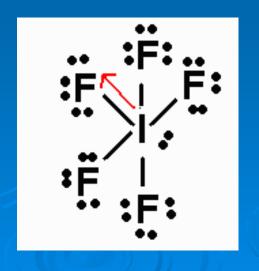


$$\dot{S} = C = \dot{S}$$



Is the following molecule polar or nonpolar: IF₅. (If polar, show the overall dipole moment.)

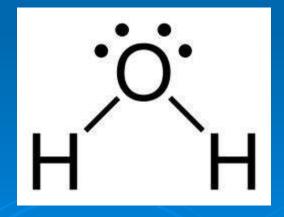
Polar – dipoles point towards the F's



- The BF₃ molecule is nonpolar, whereas the NF₃ molecule is polar. Which of the following statements accounts for the difference in polarity of the two molecules?
- a. In NF₃, each F is joined to N with multiple bonds, whereas in BF₃, each F is joined to B with single bonds.
- b. N-F bonds are polar, whereas B-F bonds are nonpolar.
- c. NF₃ is an ionic compound, whereas BF₃ is a molecular compound.
- d. Unlike BF₃, NF₃ has a nonplanar geometry due to an unshared pair of electrons on the N atom.

What is the hybridization of the molecule H₂O?

sp³



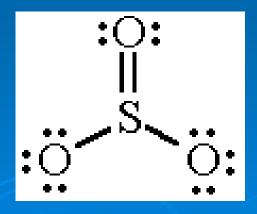
What is the hybridization of the molecule CO₂?

sp

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

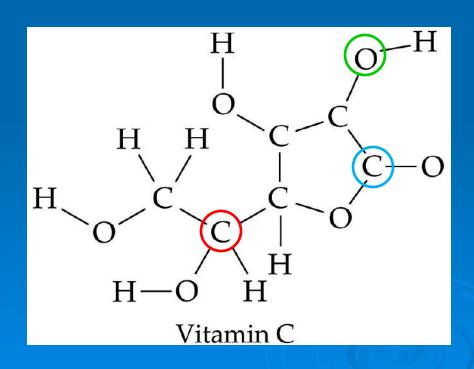
What is the hybridization of the molecule SO₃?

sp²



Below is the structure for vitamin C.

List the hybridization for each circled atom.



Green =
$$sp^3$$

Blue = sp^2
Red = sp^3

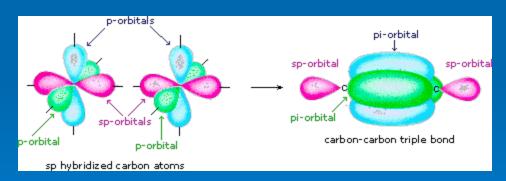
Which of the following molecules represented below contains carbon with sp² hybridization?

- a. CH₄
- b. CH₂Cl₂
- c. C_2H_6
- d. C₂H₂Cl₂
- e. C₂H₄Cl₂

d.

How many sigma and pi bonds are in N_2 ?

1 sigma and 2 pi bonds





Are the bonds in SO₃ localized or delocalized?

delocalized

$$\begin{array}{c} O \\ S=O \\ O \end{array} \begin{array}{c} O \\ S=O \end{array}$$

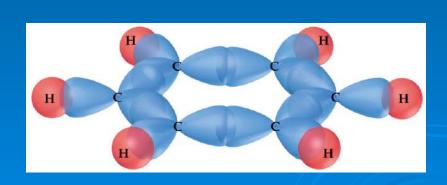
Are the bonds in NO₂- localized or delocalized?

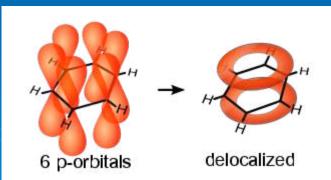
delocalized

$$-io.'_{N} \circ io. \longrightarrow io.'_{N} \circ io.'$$

Explain the sigma and pi bonds in benzene.

The C-H bonds are all sigma bonds. The C-C/C=C bonds are delocalized pi and sigma bonds.





How many sigma and pi bonds are in 1 – propene (C₃H₆)

8 sigma bonds and 1 pi bond

