# CHAPTER 14: COVALENT BONDING: ORBITALS

#### PROBLEMS WITH LEWIS STRUCTURE

- · Provides no indication of the snapes of the molecules
- · Or mould be predicted to be diamagnetic, but it is paramagnetic

Poramagnetic = has unpointed electrons.

The Levels structure of O<sub>2</sub> has an O-O double bond

with no unpaired electrons, i.e. dismogratic.

- Many compounds connot be represented by a single structure
- · Resomance structures differ only in position of electrons not the nuclei.
- · Result in notions of hupervalent compounds and "expansion of octet"
- · can't explain molecules" such as Batho Lelectron deficient molecules)

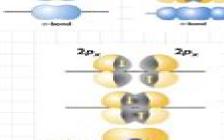
#### VALENCE—BOND THEORY "LOCALIZED ELECTRON MODEL

· Covalent bond is created by the overlap of orbitals on two atoms

#### TYPES OF OVERLAP: O and IT Bonds

cylindrically symmetrical

- · Bonding in Ha
  - sausage shaped orbital, bond called \(\sigma\)-orbital
    called \(\sigma\) orbital because it has no node and is
  - Eléction defisity is concentrated between two nuclei.
    lout is still highest at the nuclei Themselves.
- · Bonding in Na
  - · or orbital is obtained by overlap of two pr-orbitals
- Perpendicular p orbitals overlap in a side-by-side manner.
- The resulting orbital has two lobes of electron density and is called a n-bond
- called Ti-orbital because it has a node and, when viewed down the intermuclear axis, it looks like a p-orbital
- it bond has no electron density between the nuclei along the bond axis.



# <u>University Chemistry Chapter Covalent Bonding Orbitals</u> <u>Study Guide</u>

C. Barry Carter, M. Grant Norton

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