

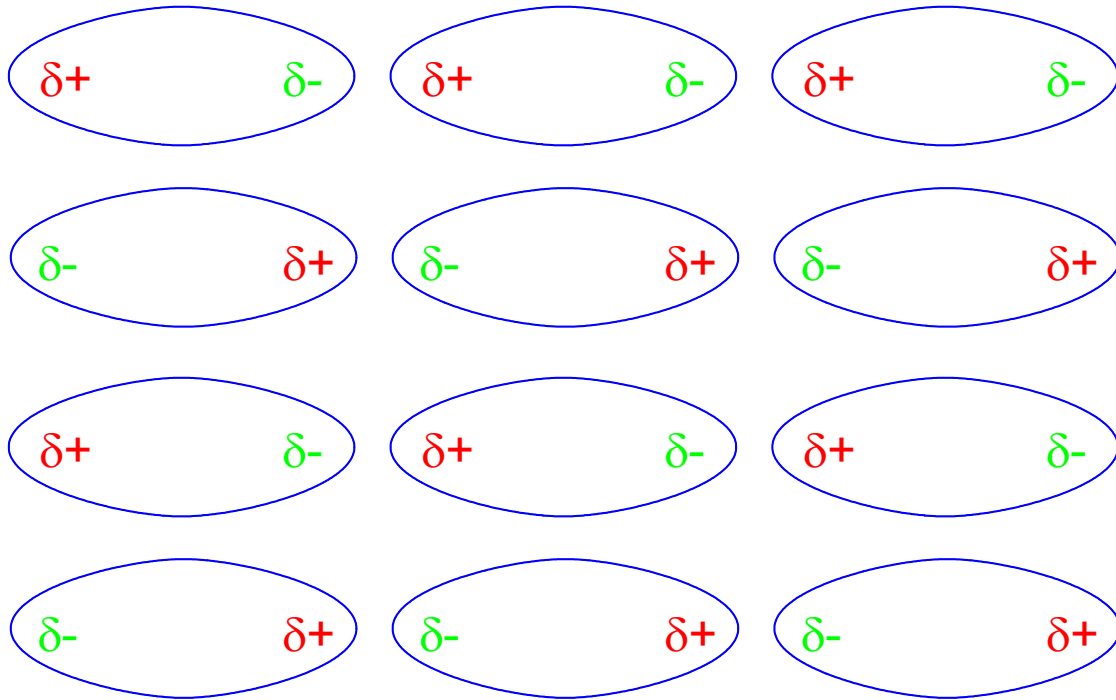
## Types of Intermolecular Forces

- van der Waals forces:
    - dipole-dipole
    - London dispersion
  - Hydrogen bonding
- ☞ Molecules can have one, two, or all three kinds of intermolecular forces, but all have London dispersion forces.
- ☞ Substances with stronger overall intermolecular forces, whatever their types, are more likely to have:
- higher melting points (m.p.)
  - higher boiling points (b.p.)
  - higher enthalpies of vaporization ( $\Delta H_{\text{vap}}$ )
  - lower vapor pressures (v.p.)
  - a condensed phase (solid or liquid) at room temperature
- ☞ Typical intermolecular forces tend to be weaker than typical covalent bond strengths.
- Covalent bonds:  $\approx 50 - 950$  kJ/mol
  - Intermolecular forces:  $\approx 1 - 50$  kJ/mol

# Dipole-Dipole Attractions Between Polar Molecules

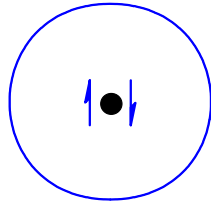
## Permanent Dipole Moments

$$D \approx 1 - 10 \text{ kJ/mol}$$

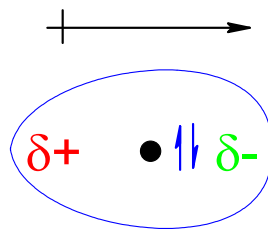


# London Dispersion Forces Transitory Dipole Moments

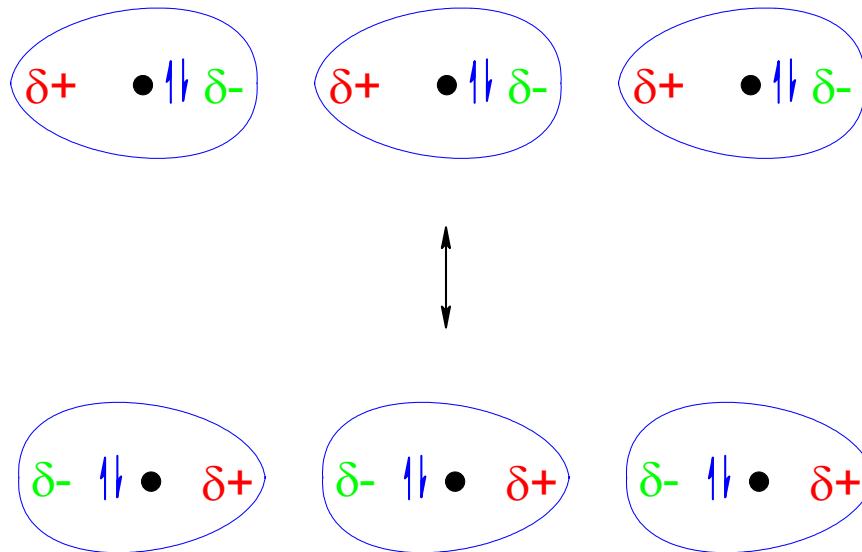
Average Electron Distribution  
No Polarity



Momentary Electron Distribution  
Temporary Polarity



## London Dispersion Forces Induced Transitory Dipole Moments



- ☞ London dispersion forces arise from changing electron distributions.
- ☞ All molecules have electrons, so all have London dispersion forces, regardless of whether they are polar or non-polar.
- ☞ **Polarizability** - Molecules with higher molecular weights have more electrons, generally more loosely held, resulting in electron distributions that are more susceptible to deformation from adjacent charges.
- ☞ **Heavier molecules have higher London dispersion forces.**

## Effects of London Dispersion Forces on Properties

### Group VI Hydrides, H<sub>2</sub>X

|           |                  |                   |                   |
|-----------|------------------|-------------------|-------------------|
| Compound  | H <sub>2</sub> S | H <sub>2</sub> Se | H <sub>2</sub> Te |
| b.p. (°C) | -60.33           | -41.3             | -2                |

### Halogens, X<sub>2</sub>

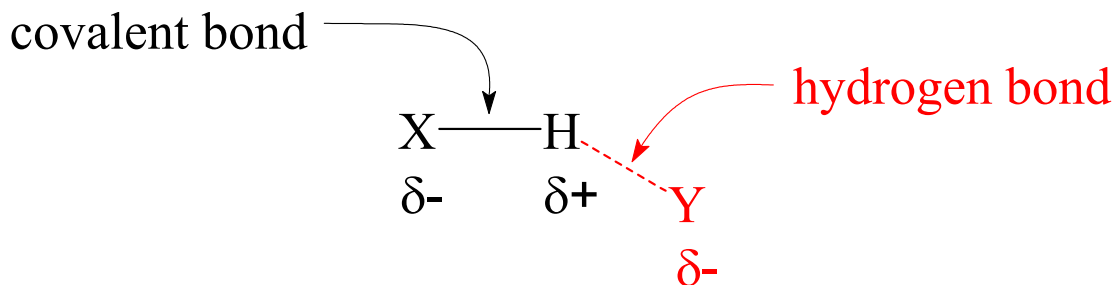
|           |                |                 |                 |                |
|-----------|----------------|-----------------|-----------------|----------------|
| Element   | F <sub>2</sub> | Cl <sub>2</sub> | Br <sub>2</sub> | I <sub>2</sub> |
| m.p. (°C) | -220           | -101            | -7.3            | 114            |
| b.p. (°C) | -188           | -34             | 58.8            | 184            |
| At 25 °C  | gas            | gas             | liquid          | solid          |

### Alkanes (C<sub>n</sub>H<sub>2n+2</sub>) at Room Temperature

| <i>n</i> | Phase  | Examples                              |
|----------|--------|---------------------------------------|
| 1 - 4    | gas    | methane, ethane, propane, butane      |
| 5 - 17   | liquid | pentane, hexane, heptane, octane, ... |
| 18 - ∞   | solid  | paraffin waxes                        |

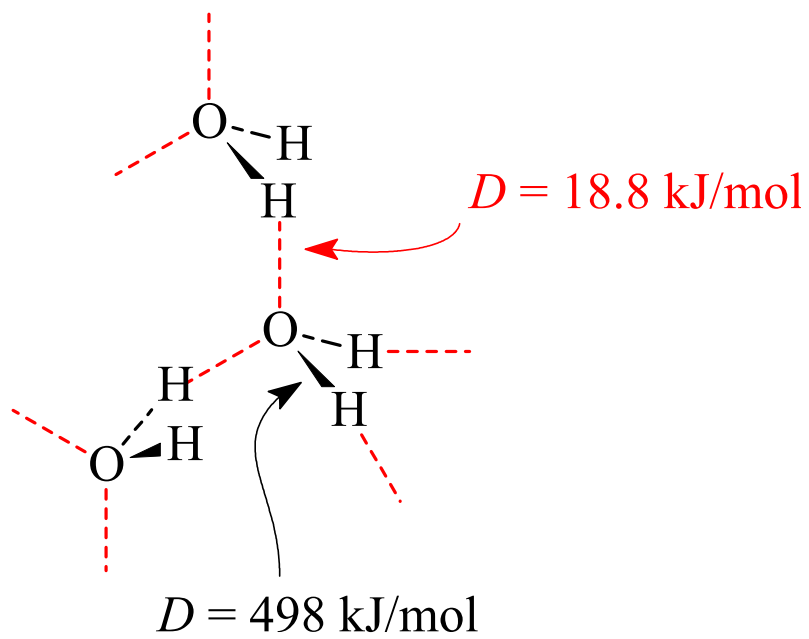
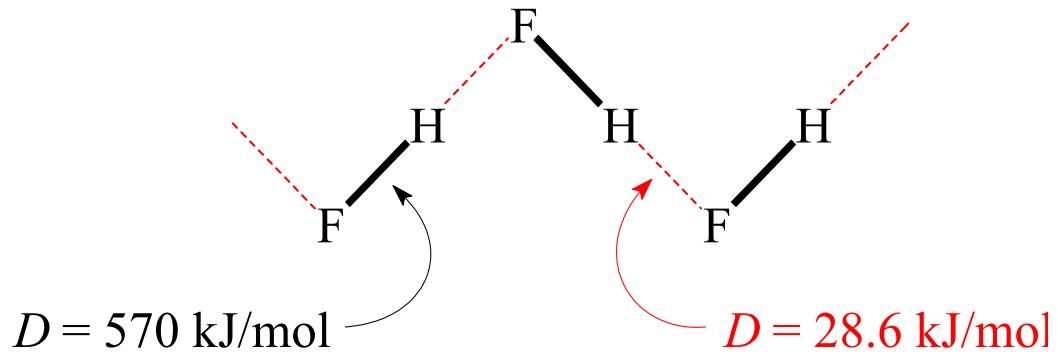
## Hydrogen Bonding

- ☞ A hydrogen bond is a non-covalent attraction between a hydrogen that is covalently bonded to a very electronegative atom (X) and another very electronegative atom (Y), most often on an adjacent molecule. (X and Y may be the same or different elements.)

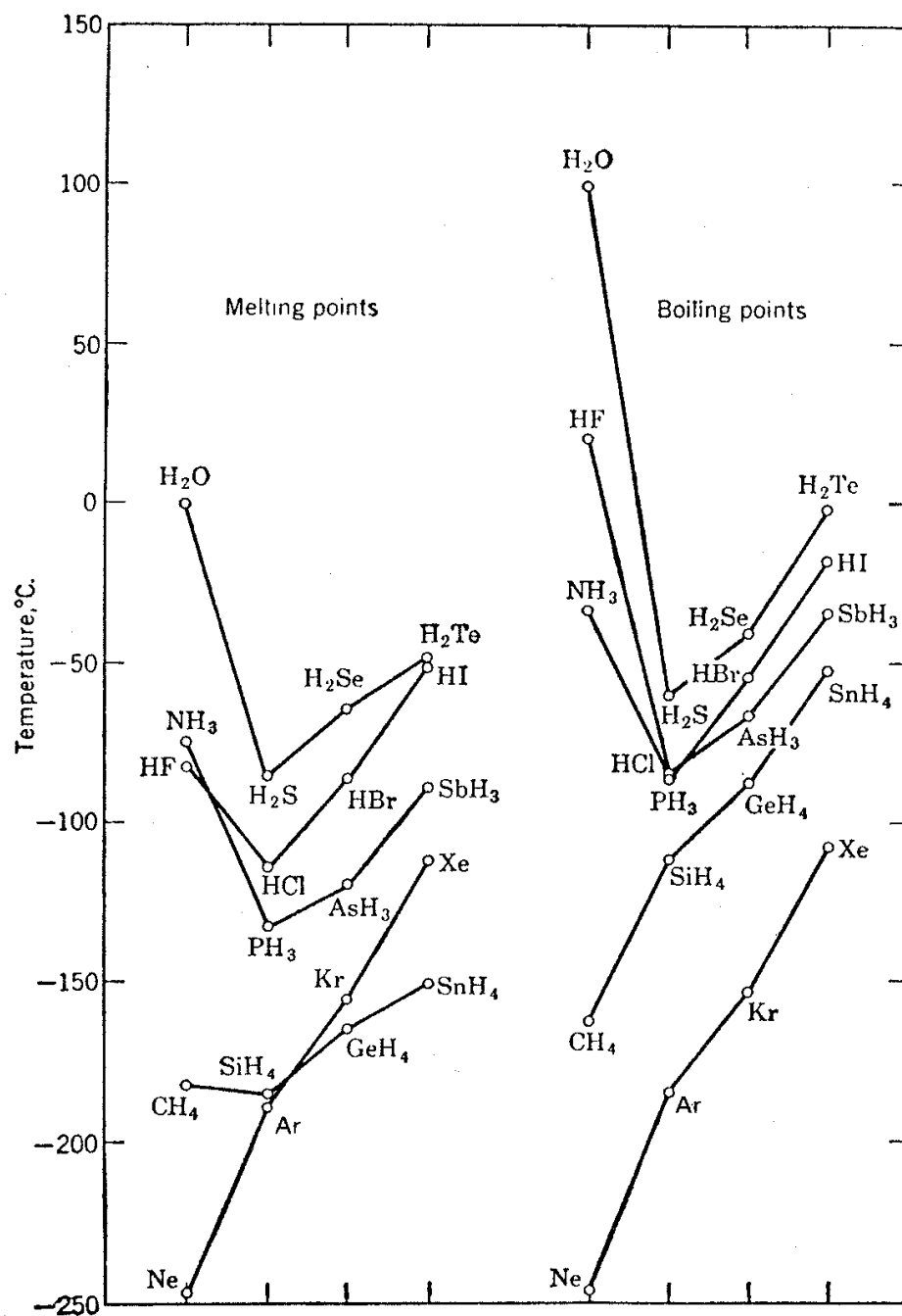


- X and Y are typically O, N, or F.
- The X–H covalent bond must be very polar.
- The hydrogen bond H $\cdots$ Y is mainly electrostatic.
- Hydrogen bond strengths typically are in the range 4 - 46 kJ/mol.

## Hydrogen Bonding in Solid HF and H<sub>2</sub>O



# Melting Points and Boiling Points of Hydrides, $XH_n$ ( $n = 1, 2, 3, 4$ )





# Structure of Ice

