

Some Background

I. Gases:

- A. \_\_\_\_\_ to fill containers
- B. Highly \_\_\_\_\_
- C. Low \_\_\_\_\_!

II. Pressure:

- A. \_\_\_\_\_
- B. Most common unit: \_\_\_\_\_
- C. Other units: \_\_\_\_\_

III. STP:

- A. \_\_\_\_\_
- B. This "standard" applies \_\_\_\_\_.

IV. Temperature:

- A. Measures \_\_\_\_\_
- B. Always use \_\_\_\_\_ for gas problems!
- C. \_\_\_\_\_.

Basic Gas Laws

I. \_\_\_\_\_ Law:

- A. Relates \_\_\_\_\_ and \_\_\_\_\_.
- B. \_\_\_\_\_ and \_\_\_\_\_ are \_\_\_\_\_.
- C. \_\_\_\_\_

II. \_\_\_\_\_ Law:

- A. Relates \_\_\_\_\_ and \_\_\_\_\_.
- B. \_\_\_\_\_ and \_\_\_\_\_ are \_\_\_\_\_.
- C. \_\_\_\_\_

III. \_\_\_\_\_ Law:

- A. Relates \_\_\_\_\_ and \_\_\_\_\_.
- B. \_\_\_\_\_ and \_\_\_\_\_ are \_\_\_\_\_.
- C. \_\_\_\_\_

IV. \_\_\_\_\_ Law:

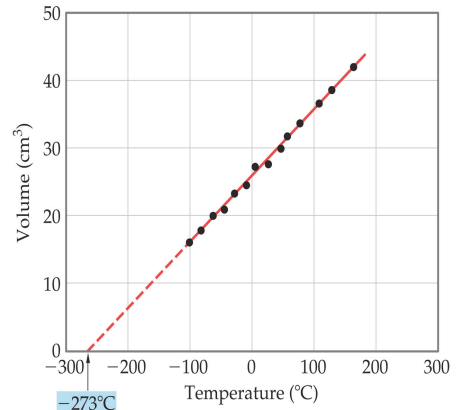
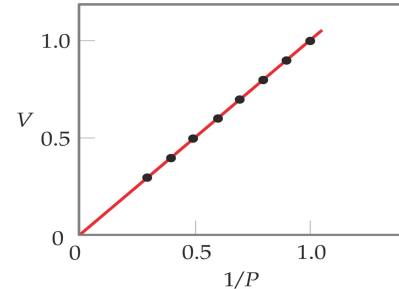
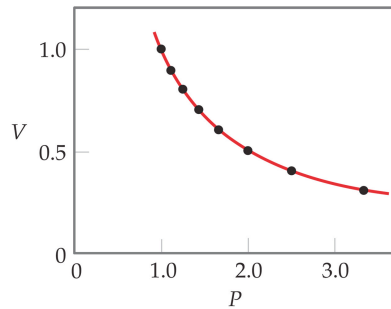
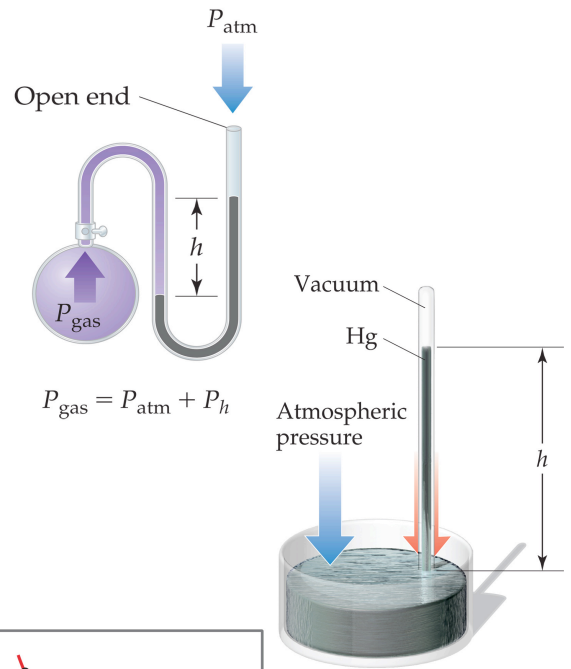
- A. Relates \_\_\_\_\_ and \_\_\_\_\_.
- B. \_\_\_\_\_.
- C. \_\_\_\_\_ and \_\_\_\_\_ are \_\_\_\_\_.
- D. \_\_\_\_\_.

E. Thus, at STP, \_\_\_\_\_ = \_\_\_\_\_. (more on this later...)

Ideal Gas Law

I. Combo of all the gas laws

- A. \_\_\_\_\_.
- B. VERY commonly used in AP for gas problems!
- C.  $R =$  \_\_\_\_\_ or \_\_\_\_\_.
- D. Derived from the \_\_\_\_\_ law:



II. Extensions of the ideal gas law:

- A. Can find density (derivation)
  
- B. Can find molecular mass (derivation)

Gas Mixtures: \_\_\_\_\_

I. Previous slides dealt with pure gases.

- A. For mixtures of gases: \_\_\_\_\_ = \_\_\_\_\_.
- B. \_\_\_\_\_
- C. When water is present, water vapor \_\_\_\_\_. (Thus, don't forget to account for it!)

II. Partial pressures can relate to moles...

- A. For mixtures of gases: gases behave \_\_\_\_\_.
- B. Therefore, we can relate \_\_\_\_\_ to \_\_\_\_\_
- C. \_\_\_\_\_ = \_\_\_\_\_ = \_\_\_\_\_.
- 1. Equation =

- D. \_\_\_\_\_ relates to \_\_\_\_\_.
- 1. Equation:

Kinetic Molecular Theory

I. This theory is essential for the gas laws to work

II. It states:

- A. Gases have \_\_\_\_\_
- B. Gases move \_\_\_\_\_
- C. Gas molecules have \_\_\_\_\_
- D. \_\_\_\_\_ between gas molecules are \_\_\_\_\_ (no \_\_\_\_\_)
- E. Average kinetic energy of molecules is \_\_\_\_\_

Graham's Law of Diffusion/Effusion

I. Introducing a unit: \_\_\_\_\_

- A. Since there are so many gas molecules in a given volume, \_\_\_\_\_

- B. Related to kinetic energy. \_\_\_\_\_
- 1. higher the \_\_\_\_\_, higher the \_\_\_\_\_.

C. Thus, \_\_\_\_\_ !

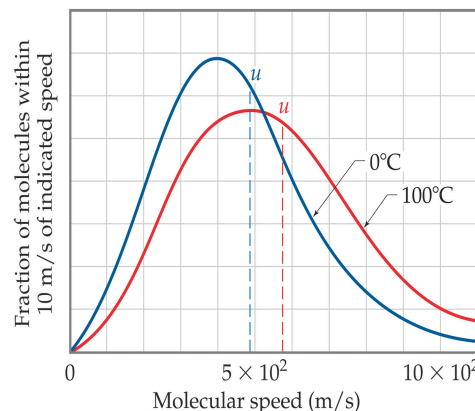
D. An equation is used to describe this:

$$u = \frac{\sqrt{3RT}}{\sqrt{M}} \quad R = \text{_____} \quad T = \text{_____}$$

$$M = \text{_____}$$

E. Thus, \_\_\_\_\_.

II. Effusion – \_\_\_\_\_.



Chapter 5: Gases

- A. Graham's law relates \_\_\_\_\_ to \_\_\_\_\_.
- B. \_\_\_\_\_.
- C. Equation: \_\_\_\_\_
- D. Diffusion: \_\_\_\_\_

Real Gases - Van der Waals Equation

- I. Not all gases are \_\_\_\_\_
- II. The closer the gas is to \_\_\_\_\_, the \_\_\_\_\_  
(\_\_\_\_\_).
- III. Ideal gas law makes two assumptions that \_\_\_\_\_:
1. \_\_\_\_\_
  2. \_\_\_\_\_
- IV. van der Waals equation accounts for this.

1.  $a =$  \_\_\_\_\_ (units: \_\_\_\_\_)
2.  $b =$  \_\_\_\_\_ (units: \_\_\_\_\_)

Substance	$a$ (L <sup>2</sup> -atm/mol <sup>2</sup> )	$b$ (L/mol)
He	0.0341	0.02370
Ne	0.211	0.0171
Ar	1.34	0.0322
Kr	2.32	0.0398
Xe	4.19	0.0510
H <sub>2</sub>	0.244	0.0266
N <sub>2</sub>	1.39	0.0391
O <sub>2</sub>	1.36	0.0318
Cl <sub>2</sub>	6.49	0.0562
H <sub>2</sub> O	5.46	0.0305
CH <sub>4</sub>	2.25	0.0428
CO <sub>2</sub>	3.59	0.0427
CCl <sub>4</sub>	20.4	0.1383

V. Thus...

- A. The more \_\_\_\_\_ a molecule is, the \_\_\_\_\_ it is.
1. The \_\_\_\_\_ affects its \_\_\_\_\_.
  2. Polarizable: \_\_\_\_\_
- B. The \_\_\_\_\_
1. \_\_\_\_\_, causing \_\_\_\_\_ to come into play.
  2. \_\_\_\_\_, \_\_\_\_\_ for the molecules to be \_\_\_\_\_.