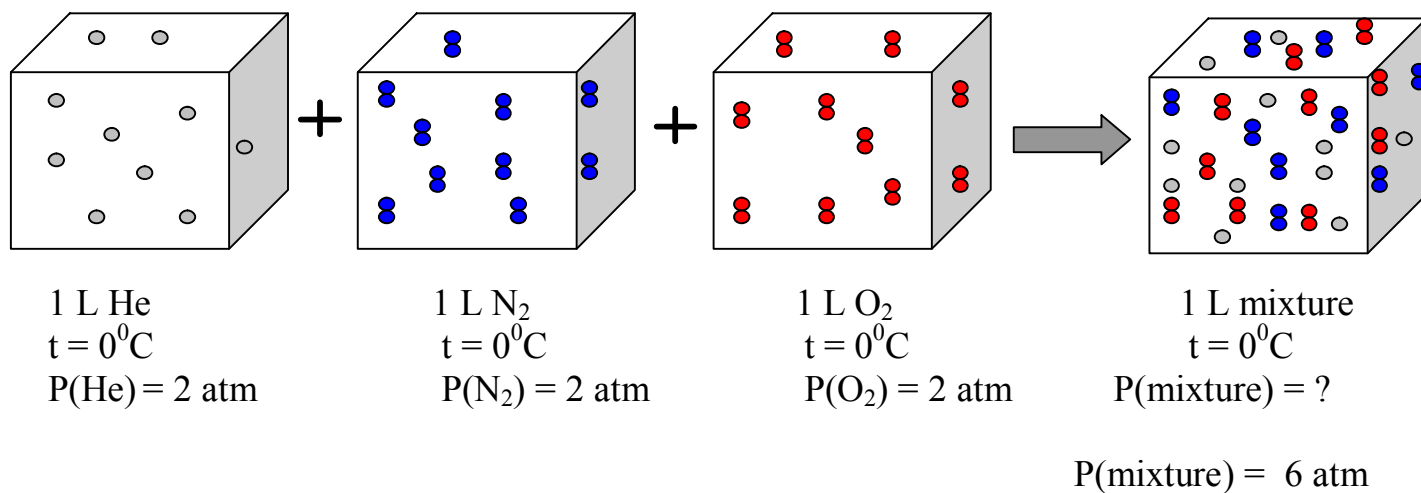


GAS MIXTURES

- Consider mixing equal volumes of 3 different gases, all at the same temperature and pressure in a container of the same size.



NOTE:

- Each gas in a mixture of gases acts as though it were the only gas in the mixture

$$P(\text{mixture}) = \begin{matrix} P(\text{He}) & + & P(\text{N}_2) & + & P(\text{O}_2) \\ \downarrow & & \downarrow & & \downarrow \\ \text{PARTIAL PRESSURES} & & & & \end{matrix}$$

DALTON'S LAW OF PARTIAL PRESSURES

- **The Total Pressure (P_T) of a mixture of gases is equal to the sum of the partial pressures of the gases in the mixture.**

$$P_T = P_A + P_B + P_C + \dots\dots$$

MOLE FRACTION OF A GAS IN A MIXTURE OF GASES

$$\begin{array}{l} \text{Gas A + Gas B + Gas C} \\ \text{Total Number of Moles of gas in the Mixture} \end{array} = \begin{array}{l} \text{Mixture of gases} \\ N \end{array}$$

N_A = number of moles of gas A

N_B = number of moles of gas B

N_C = number of moles of gas C

$$N = N_A + N_B + N_C$$

$$\frac{N_A}{N} = \text{Mole Fraction of Gas A} = X_A$$

$$\frac{N_B}{N} = \text{Mole Fraction of Gas B} = X_B$$

$$\frac{N_C}{N} = \text{Mole Fraction of Gas C} = X_C$$

Mole Percent = Mole fraction X 100

$$\text{Example: Mole \% of A} = \frac{N_A}{N} \times 100$$

DALTON'S LAW OF PARTIAL PRESSURES

- The **Pressure** of a sample gas is **directly proportional** to the number of molecules (expressed as **number of moles**) of gas.

$$\frac{P_A}{P} = \frac{N_A}{N}$$

$$\frac{P_B}{P} = \frac{N_B}{N}$$

$$\frac{P_C}{P} = \frac{N_C}{N}$$

Where:

- N_A, N_B, N_C = number of moles of gas A, B and C respectively
- N = total number of moles of gas mixture
- P_A, P_B, P_C = partial pressures of gas A, B and C respectively
- P = total pressure of gas mixture

Alternately,

$$P_A = X_A P \quad P_B = X_B P \quad P_C = X_C P$$

Examples:

- Calculate the total pressure (in atm) and partial pressure of each gas in a mixture composed of 0.0200 moles helium gas and 0.0100 moles hydrogen gas in a 5.00 L flask at 10⁰ C.

$$N_T = 0.0200 \text{ mol} + 0.0100 \text{ mol} = 0.0300 \text{ mol}$$

$$\begin{aligned} n_T &= 0.0300 \text{ mol} \\ V &= 5.00 \text{ L} \\ T &= 283 \text{ K} \\ P &= ? \end{aligned}$$

$$P V = n_T R T \quad P = \frac{n_T R T}{V}$$

$$P = \frac{(0.300 \text{ mol})(0.0821 \frac{\text{L atm}}{\text{mol K}})(283 \text{ K})}{5.00 \text{ L}} = 0.139 \text{ atm}$$

$$P_{\text{He}} = X_{\text{He}} P =$$

$$P_{\text{H}_2} = X_{\text{H}_2} P =$$

2. The atmosphere in a sealed diving bell contains oxygen and helium. If the gas mixture has 0.200 atmospheres of oxygen and a total pressure of 3.00 atmospheres, calculate the mass of helium in 1.00 L of gas mixture at 20°C.

PART A: Calculate the pressure of helium

$$P_T = P_{\text{Oxygen}} + P_{\text{Helium}}$$

$$P_{\text{Helium}} = P_T - P_{\text{Oxygen}}$$

$$P_{\text{Helium}} = 3.00 \text{ atm} - 0.200 \text{ atm} = 2.80 \text{ atm}$$

PART B: Calculate the number of moles of Helium

$$P_{\text{He}} = 2.80 \text{ atm}$$

$$V_{\text{He}} = 1.00 \text{ L}$$

$$T_{\text{He}} = 293 \text{ K}$$

$$n_{\text{He}} = ?$$

$$PV = nRT$$

$$n = \frac{PV}{RT}$$

$$n = \frac{(2.80 \text{ atm})(1.00 \text{ L})}{(0.0821 \frac{\text{L atm}}{\text{mol K}})(293 \text{ K})} = 0.1165 \text{ mol}$$

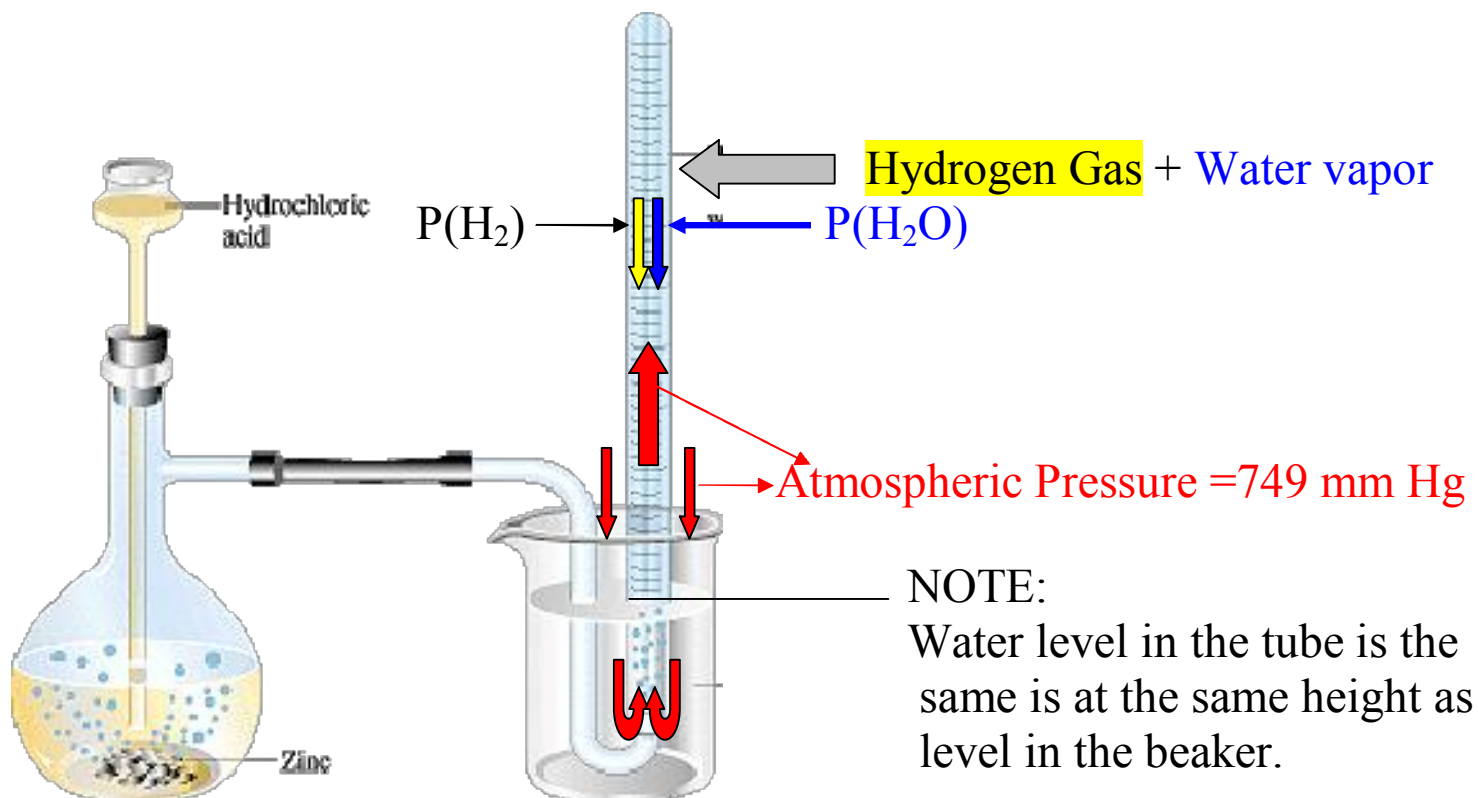
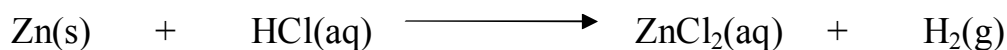
PART C: Calculate the mass of Helium

$$? \text{ g He} = 0.1165 \text{ moles He} \times \frac{4.00 \text{ g He}}{1 \text{ mole He}} = 0.466 \text{ g He}$$

3. A mixture of 0.25 mol H₂ and 0.45 mol of N₂ are placed in a 1.50-L flask at 27°C. Calculate the partial pressures of each gas and the total pressure.
4. The partial pressure of CH₄ gas is 0.175 atm and that of O₂ gas is 0.250 atm in a mixture of the two gases. What is the mole fraction of each gas in the mixture?

COLLECTING GASES OVER WATER

- Many gases are commonly collected over water.
- Consider the collection of Hydrogen gas from the reaction of zinc with hydrochloric acid:



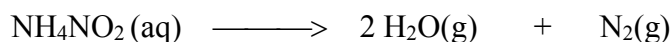
$$\begin{aligned} \text{Gas Pressure in the tube} &= P_T = P(\text{H}_2) + P(\text{H}_2\text{O}) \\ P(\text{atm}) &= P_T = P(\text{H}_2) + P(\text{H}_2\text{O}) \end{aligned}$$

$$P(\text{atm}) = P(\text{H}_2) + P(\text{H}_2\text{O})$$

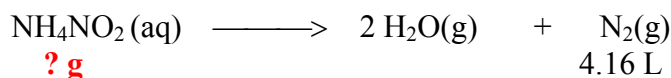
$P(\text{H}_2\text{O})$ = is temperature dependent
 = is given in tables (Table 5.6, page 206)

Example:

1. An aqueous solution of ammonium nitrite decomposes when heated to give off nitrogen gas and water vapor:



How many grams of NH_4NO_2 must have reacted if 4.16 L of N_2 gas was collected over water at 19°C and 0.965 atmospheres of pressure ?

**PART A: Calculate the pressure of dry Nitrogen**

$$P_T = P(\text{N}_2) + \text{Pressure of Water Vapor}$$

$$P(\text{N}_2) = P_T - \text{Pressure of Water Vapor}$$

$$P_T = 0.965 \text{ atm}$$

$$\text{Pressure of Water Vapor} = ?$$

$$\text{Table 5.6 at } 19^\circ\text{C} \Rightarrow P(\text{H}_2\text{O}) = 16.5\text{mmHg}$$

$$16.5 \text{ mm Hg} \times \frac{1 \text{ atm}}{760.0 \text{ mm Hg}} = 0.0217 \text{ atm}$$

$$P(\text{N}_2) = 0.965 \text{ atm} - 0.0217 \text{ atm} = \mathbf{0.9433 \text{ atm}}$$

PART B: Calculate the number of moles of nitrogen

$$P = 0.9433 \text{ atm}$$

$$V = 4.16 \text{ L}$$

$$T = 292 \text{ K}$$

$$n = ?$$

$$P V = n R T \quad n = \frac{P V}{R T}$$

$$n = \frac{(0.9433 \text{ atm})(4.16 \text{ L})}{(0.0821 \frac{\text{L} \cdot \text{atm}}{\text{mol} \cdot \text{K}})(292 \text{ K})} = \mathbf{0.1637 \text{ mol}}$$

PART C: Calculate the mass of NH_4NO_2 (Stoichiometry)

$$?g \text{ NH}_4\text{NO}_2 = 0.1637 \text{ moles N}_2 \times \frac{1 \text{ mole NH}_4\text{NO}_2}{1 \text{ mole N}_2} \times \frac{64.06 \text{ g NH}_4\text{NO}_2}{1 \text{ mole NH}_4\text{NO}_2} = \mathbf{10.5 \text{ g}}$$

- Helium is collected over water at 25°C and 1.00 atm total pressure. What volume of gas must be collected to obtain 0.586 g of helium? (At 25°C the vapor pressure of water is 23.8 mmHg)

KINETIC - MOLECULAR THEORY OF IDEAL GASES

(Theory of Gas Molecules in Motion)

 Ideal Gas = A gas that follows the IDEAL GAS LAW (**P V = n R T**)

BASIC IDEA:

- **A GAS CONSISTS OF MOLECULES IN CONSTANT RANDOM MOTION**

Recall:	$\text{Kinetic Energy} = E_K = \frac{1}{2} m v^2$	m = mass of molecule v = molecular speed E _K = energy of moving molecule
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POSTULATE 1:

- The size of molecules is negligible compared to the intermolecular distances

Consequence:

- **The Volume of a gas is determined by the distances between the molecules.**

POSTULATE 2:

- Molecules move randomly in straight lines in all directions and at various speeds

Consequence:

- **The Pressure of a gas is the same in all directions**

POSTULATE 3:

- The attractive or repulsive forces between molecules are very weak (negligible)

POSTULATE 4:

- When molecules collide with one another, the collisions are elastic (no energy is lost). This is similar to collisions between billiard balls.

POSTULATE 5:

- The Average Kinetic Energy is directly proportional to the Absolute Temperature

INTERPRETATION OF THE KINETIC-MOLECULAR THEORY OF GASES

The **PRESSURE** of a gas results from:

-The **COLLISIONS** of the molecules with the walls of the container

The NUMBER OF COLLISIONS is determined by:

-the number of molecules per unit volume
-average speed of molecules

BOYLE'S LAW

Recall: The Temperature is constant; therefore:

$$KE_{ave} = \text{constant}$$

**An Increase
In Volume**



Decreases the number of molecules per unit volume



Decreases the frequency of collisions



**Decreases the
Pressure**

CHARLES' LAW

Recall: The pressure is constant, therefore: Frequency and force of molecular collisions are constant.

**An Increase
In Temperature**



Increases Average Molecular Speed



Increases the Volume
(molecules spread), in order to maintain frequency of collisions unchanged

MOLECULAR SPEEDS

- In gases, molecular speeds vary over a wide range
- Meaning: Different molecules move at different speeds, that are constantly changing, as the molecules collide with one another and the walls of the container

It is more convenient to refer to :

U = AVERAGE MOLECULAR SPEED
= THE SPEED OF A MOLECULE HAVING AVERAGE
KINETIC ENERGY (KE_{AVE})

$$U = \sqrt{\frac{3RT}{M_m}}$$

R = gas constant

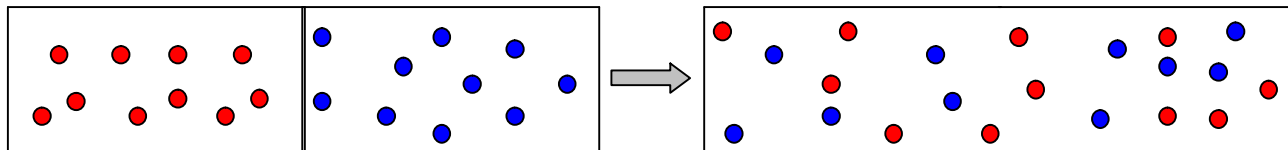
T = absolute temperature

M_m = molar mass

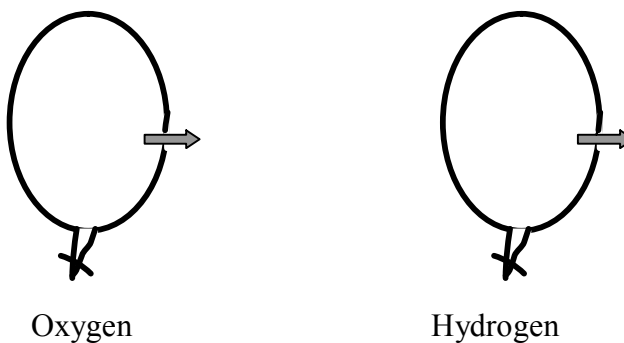
NOTE: **The higher the molar mass, the lower the average molecular speed.**
(The heavier the molecules, the slower they move)

DIFFUSION & EFFUSION

- Diffusion is the process by which a gas spreads through another gas.



- Effusion is the process by which a gas flows out from a container through a small hole.



- Same Volume, Same Pressure, Same Temperature, Same Size Hole
- Which Balloon deflates faster? OR Which gas has a **higher Rate of Effusion** ?
- What is the **Ratio between the Rates of Effusion** ?

Rate of Effusion depends on:

1. The cross-sectional area of the hole → (the same)
2. The number of molecules per unit volume → (the same)
3. **The average Molecular Speed (U)** → **(different)**
(the faster they move, the sooner they escape)

Recall:
$$U = \sqrt{\frac{3RT}{M_m}}$$

NOTE: The Rate of Effusion:

- is directly proportional to the Average Molecular Speed (U)
- is inversely proportional to the square root of the Molar Mass

GRAHAM'S LAW OF EFFUSION

- **The Rate of Effusion is inversely proportional to the square root of the Molar Mass**

$$\frac{\text{Rate of Effusion of H}_2}{\text{Rate of Effusion of O}_2} = \frac{\sqrt{\text{Molar Mass of O}_2}}{\sqrt{\text{Molar Mass of H}_2}} = \sqrt{\frac{32.0}{2.0}} = 4$$

Conclusion: Hydrogen will effuse 4 times faster than Oxygen

Examples:

1. Obtain the ratio of effusion of H₂ and H₂S under the same conditions.

$$\frac{\text{Rate H}_2}{\text{Rate H}_2\text{S}} = \frac{\sqrt{M_m(\text{H}_2\text{S})}}{\sqrt{M_m(\text{H}_2)}} = \sqrt{\frac{34}{2}} = 4.1$$

2. If 0.10 mol of I₂ vapor can effuse from an opening in a heated vessel in 52 seconds, how long will it take 0.10 mol of H₂ to effuse under the same condition?

$$\frac{\text{Rate H}_2}{\text{Rate I}_2} = \frac{\sqrt{M_m(\text{I}_2)}}{\sqrt{M_m(\text{H}_2)}} = \sqrt{\frac{253.8}{2.02}} = 11.2$$

H₂ effuses 11.2 faster than I₂

It follows that the time t takes H₂ to effuse is 11.2 times shorter:

$$T(\text{H}_2) = \frac{52 \text{ seconds}}{11.2} = 4.6 \text{ seconds}$$

REAL GASES

Gases behave according to the Ideal Gas Law, provided:

1. The space occupied by molecules is truly negligible compared to the total gas volume (Postulate 1),
2. The intermolecular forces of attraction are truly negligible (Postulate 3)

THIS IS TRUE:

- at High Temperature, and
- at Low Pressure

Reasons:

- Molecules are far apart
- Molecules do not attract

IDEAL GAS

$$PV = nRT$$

THIS IS NOT QUITE TRUE:

- at Low Temperature, and
- at High Pressure

Reasons:

- Molecules are relatively closer

As such:

- **Volume occupied by molecules must be taken into account**
- **Intermolecular forces between molecules are not negligible and molecules begin to attract one another**

$$\left(P + \frac{n^2 a}{V^2}\right)(V - nb) = nRT$$

Van Der Waals Equation for Real Gases

NOTE:

V (intermolecular space) becomes **(V - nb)**
 - **decreased Volume** accounts for space occupied by molecules)

P becomes $\left(P + \frac{n^2 a}{V^2}\right)$ **increased Pressure** accounts for intermolecular forces of attraction

a and **b**- are constants
 - have different values for different gases