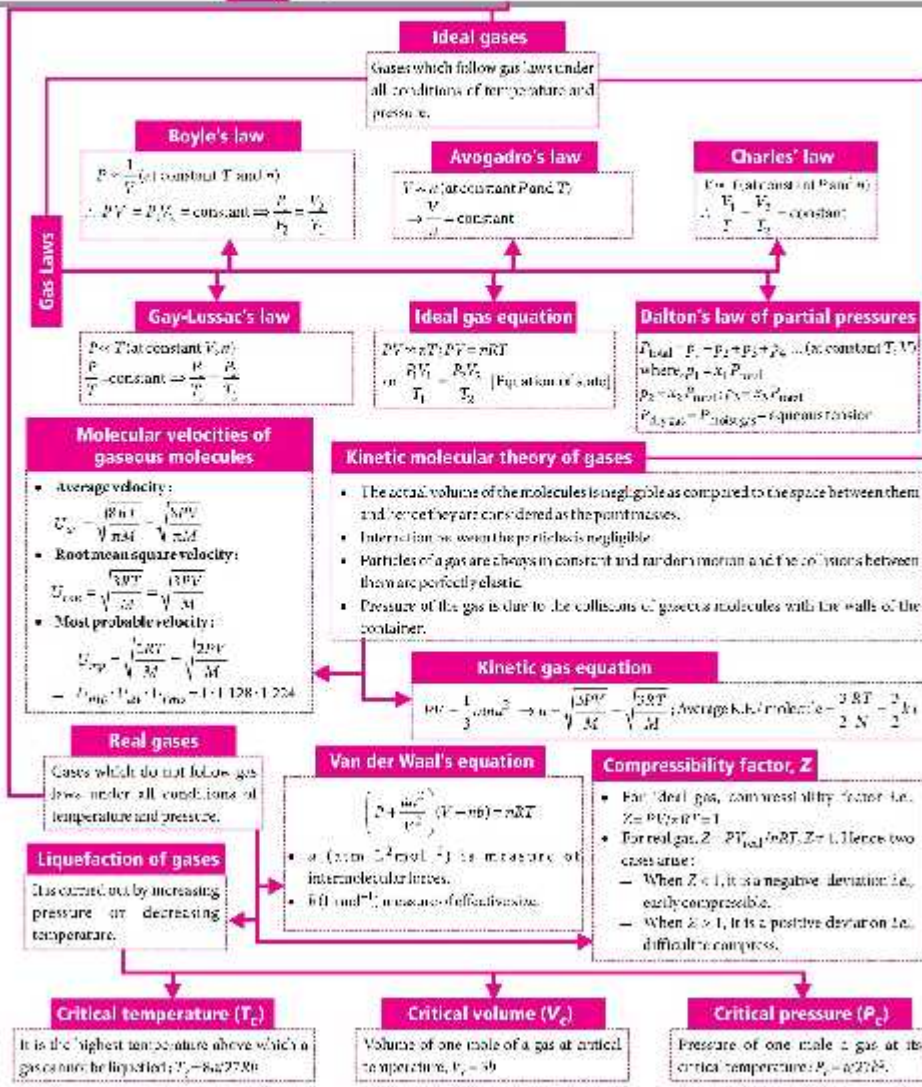


CONCEPT MAP

GASEOUS STATE

A substance exists in gaseous state if the thermal energy of molecules predominates over the attractive forces, molecules are sufficiently apart from one another (intermolecular distances $10^{-7} - 10^{-8}$ cm), have large constant vibratory and translatory motions and are most energetic.

Class XI



CONCEPT MAP CLASS XI

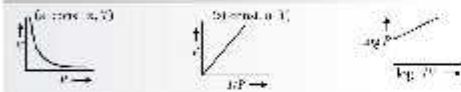
Gaseous State

A gas occupies volume about 1600 times greater than that of the equal weight of liquid. If gas is compressed to 1/1600th of its volume above critical temperature (which is different for each gas) no phase change occurs and the resulting substance is a gas that is just as dense as a liquid.

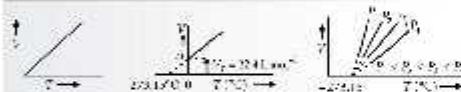
Gas Laws

- Boyle's law**: It states that at a constant temperature (T), $P \propto 1/V$ or $PV = \text{constant}$ or $P_1V_1 = P_2V_2$.
- Charles' law**: It states that at constant pressure (P), $V \propto T$ or $(V/T) = \text{constant}$ or $V_1/T_1 = V_2/T_2$.
- Avogadro law**: It states that at constant temperature (T) and pressure (P), $V \propto n$ or $(V/n) = \text{constant}$.
- Gay Lussac's law**: It states that at constant volume (V), $P \propto T$ or $(P/T) = \text{constant}$.

Boyle's Law



Charles' Law



Ideal Gas Equation

$PV = nRT$ (R = gas constant)

$0.0821 \text{ L atm K}^{-1} \text{ mol}^{-1} = \frac{R}{8.314 \text{ J K}^{-1} \text{ mol}^{-1}} = \frac{R}{\text{gas constant}} = 0.083 \text{ L bar K}^{-1} \text{ mol}^{-1} = 1.99 \text{ cal K}^{-1} \text{ mol}^{-1}$

Deviation from Ideal Gas Behaviour (Real Gas)

- Real gas follows gas laws only at high temperature and low pressure.
- It follows van der Waal's equation: $\left(P + \frac{a}{V^2}\right)(V - nb) = nRT$

Compressibility Factor

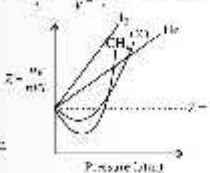
$Z = \frac{PV}{nRT}$

• For ideal gas, compressibility factor i.e., $Z = PV/nRT = 1$

• For real gas, $Z = PV/nRT$, $Z \neq 1$. Hence two cases arise:

— When $Z < 1$, it is a negative deviation i.e., easily compressible.

— When $Z > 1$, it is a positive deviation i.e., difficult to compress.



- Relation between critical constants and van der Waal's constants:** $V_c = 3b$, $P_c = \frac{a}{27b^2}$, $T_c = \frac{8a}{27Rb}$

- As the value of 'a' increases, the critical temperature (T_c) also increases. Higher the critical temperature of the gas, more easily it liquefies.
- The ease of liquefaction order: $\text{Cl}_2 > \text{CO}_2 > \text{CH}_4 > \text{H}_2 > \text{He}$

Graham's Law of Diffusion

$$\frac{r_1}{r_2} = \frac{v_1/t_1}{v_2/t_2} = \sqrt{\frac{M_2}{M_1}} = \sqrt{\frac{d_2}{d_1}}$$

Dalton's Law of Partial Pressure

It states that "at a given temperature, the total pressure (P_{total}) exerted by two or more non reacting gases occupying a definite volume is equal to the sum of partial pressures of the component gases".

$$P_{\text{total}} = P_1 + P_2 + P_3 + \dots \text{ (where } P_1, P_2, P_3 \text{ are partial pressures of component gases)}$$

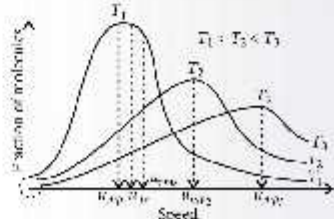
Kinetic Molecular Theory of Gases

- All gases are made up of a very large number of minute particles like atoms or molecules.
- The molecules are in continuous random motion in all directions.
- Molecular collisions are perfectly elastic.
- The intermolecular forces of attraction are negligible.
- The average kinetic energy of a gas molecule is directly proportional to the absolute temperature.
- Kinetic gas equation: $PV = \frac{1}{3} nma^2$

Molecular Speed

- Root mean square velocity: $u_{\text{rms}} = \sqrt{\frac{3RT}{M}}$
- Average velocity: $u_{\text{av}} = \sqrt{\frac{8RT}{\pi M}}$
- Most probable speed: $u_{\text{mp}} = \sqrt{\frac{2RT}{M}}$
- $u_{\text{mp}} : u_{\text{av}} : u_{\text{rms}} = 1 : 1.128 : 1.224$

Maxwell-Boltzmann Distribution Curve



Signifies that speed is never zero but it tends towards zero.