		···· The instation of		(a) 1.5 <i>lit.</i> (c) 11.2 <i>lit.</i>	 (b) 2.8 <i>lit.</i> (d) 22.4 <i>lit.</i>
	🚽 Ordina	iry i ninking	12.	Pressure of a gas in a vesse	l can be measured by
		, ,		(a) Barometer	(b) Manometer
		Objective Questions		(c) Stalgometer	(d) All the baove
Cha	aracteristics and Measu	rable properties of gases	13.	Volume occupied by a gas <i>V mL.</i> Its volume at 273 <i>K</i>	at one atmospheric pressure and $0^{o}C$ is will be
	Which and of the following state	mente is not compatible that the three			[Bihar MADT 1982]
•	states of matter <i>i.e.</i> solid, liquid	and gaseous		(a) V ml	(b) $V/2 ml$
	(a) Molecules of a solid posse	ss least energy whereas those of a		(c) $2V$	(d) None of these
	gas possess highest energy	/	14.	Which one of the following	statements is wrong for gases
	(b) The density of solid is high	hest whereas that of gases is lowest		(a) Gases do not have a	definite shape and volume
	(c) Gases like liquids possess	definite volumes		(b) Volume of the gas is	equal to the volume of the container
	(d) Molecules of a solid posses	ss vibratory motion		confining the gas	
2.	The temperature and pressure a vapour can exist together are	at which ice, liquid water and water		(c) Confined gas exerts u container in all direct	iniform pressure on the walls of its tions
	(a) $0^{o}C, 1 atm$	(b) $2^{o}C, 4.7 atm$		(d) Mass of the gas cannot in which it is enclosed	ot be determined by weighing a container d
	(c) $0^{o} C, 4.7 mm$	(d) $-2^{o}C, 4.7 mm$	15.	Which of the following exhi	ibits the weakest intermolecular forces [AIIMS
.	Which of the following is true a	bout gaseous state		(a) NH_3	(b) <i>HCl</i>
	(a) Thermal energy = Molecul	ar attraction		(c) He	(d) H_2O
	(b) Thermal energy >> Molecu	llar attraction	16	No is found in a litra f	ask under $100kPa$ pressure and Ω is
	(c) Thermal energy << Molecu	lar attraction	10.		$1 1 220 k \text{P}_{2}$
	(d) Molecular forces >> Those	in liquids		found in another 3 litre fia	ask under 520 KPa pressure. If the two
.	Kinetic energy of molecules is hi	ghest in		flasks are connected, the re	Suitant pressures is
	(a) Gases	(b) Solids		(a) 310 kPa	(b) 210 kPa
	(c) Liquids	(d) Solutions		(c) 420 kPa	(d) $365 \ kPa$
	Which of the following statement	nt is correct		(e) $265 \ kPa$	(1) 000 1
	(a) In all the three states the	molecules possess random	_		
	translational motion	1		Ideal gas equation	n and Related gas laws
	(b) Gases cannot be converted liquid state	i into solids without passing through		If R 1/ Transcont process	we waken and temperature of the sec
	(c) One of the common prop	erty of liquids and gases is viscosity	١.	the correct represent press	of Boyle's law is
	(d) According to Boyle's law 1	//P is constant at constant T		· ·	[BIT Ranchi 1988]
.	A volume of 1 m^3 is equal to			(a) $V \propto \frac{1}{2}$ (at constant)	\vec{p} (b) $\vec{P}\vec{V} = \vec{P}T$
	() 1000 3	(1) 100 3		(a) $V \propto \frac{1}{T}$ (at constant	F = (0) I V = KI
	(a) $1000 cm^2$	(b) $100 cm^2$		(c) $V \propto 1/P$ (at consta	nt T (d) PV = nRT
	(c) $10 dm^3$	(d) $10^6 cm^3$	2	At constant temperature in	a given mass of an ideal gas
,	Which one of the following is no	at a unit of pressure	2.	ne constante temperature, n	[CRSE PMT 1991]
•	(a) Newton	(b) Torr		(a) The ratio of pressure	and volume always remains constant
	(c) Pascal	(d) Bar		(b) Volume always remain	ns constant
				(c) Pressure always remai	ns constant
5.	$1^{\circ}C$ rise in temperature is equ	al to a rise of		(d) The product of pressu	ire and volume always remains constant
	(a) $1^{o} F$	(b) $9/5^{o}F$	3.	Air at sea level is dense. Th	is is a practical application of
	$() = 5 / 0^{9} E$	(1) $22^{0} E$	0.		[Kerala CEE 2000]
_	$(c) \Im / \Im F$	(d) 33 F		(a) Boyle's law	(b) Charle's law
) .	Which of the following relations	s for expressing volume of a sample		(c) Avogadro's law	(d) Dalton's law
	(a) $1L = 10^3 ml$	(b) $1 dm^3 = 1 L$	4.	If 20 cm^3 gas at 1 <i>atm.</i> is	expanded to 50 cm^3 at constant <i>T</i> , then
	(a) $1L - 10^3 m^3$	(4) $1I - 10^3 \text{ cm}^3$		what is the final pressure	[CPMT 1988]
	(c) 1 L = 10 m	(a) 1L = 10 cm		(a) $20 \times \frac{1}{50}$	(b) $50 \times \frac{1}{20}$
).	One atmosphere is numerically e	equal to approximately		50	20
	(a) $10^6 \text{dynes} cm^{-2}$	(b) $10^2 \text{dynes} cm^{-2}$		(c) $1 \times \frac{1}{2} \times 50$	(d) None of these
	(c) $10^4 \text{dvnes} cm^{-2}$	(d) $10^8 \text{dvnes} cm^{-2}$		20	
		-	5.	Which of the following stat	ement is false [BHU 1994]
•	$2gm$ of O_2 at 27^oC and 760	mm of Hg pressure has volume[BCECE	2005]	(a) The product of press is independent of tem	ure and volume of fixed amount of a gas perature

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- (b) Molecules of different gases have the same K.E. at a given temperature
- The gas equation is not valid at high pressure and low (c) temperature
- (d) The gas constant per molecule is known as Boltzmann constant
- 6. Which of the following graphs represent Boyle's law



- Densities of two gases are in the ratio 1 : 2 and their temperatures 7. are in the ratio 2 : 1, then the ratio of their respective pressures is [BHU 2000]
 - (b) 1:2 (a) 1:1
 - (c) 2:1 (d) 4:1
- At constant pressure, the volume of fixed mass of an ideal gas is 8. [EAMCET 1985] directly proportional to
 - (a) Absolute temperature (b) Degree centigrade
 - (c) Degree Fahrenheit (d) None
- Which of the following expression at constant pressure represents 9. Charle's law [AFMC 1990]
 - (a) $V \propto \frac{1}{T}$ (b) $V \propto \frac{1}{T^2}$
 - (d) $V \propto d$ (c) $V \propto T$
- Use of hot air balloons in sports and meteorological obsevations is 10. an application of [Kerala MEE 2002]
 - (a) Boyle's law (b) Newtonic law (c) Kelvin's law (d) Charle's law
- 11. A 10 g of a gas at atmospheric pressure is cooled from $273^{\circ}C$ to

 $0^{o}C$ keeping the volume constant, its pressure would become

(a)	1/2 <i>atm</i>	(b)	1/273 <i>atm</i>
(c)	2 <i>atm</i>	(d)	273 <i>atm</i>

Pressure remaining the same, the volume of a given mass of an ideal 12. gas increases for every degree centigrade rise in temperature by definite fraction of its volume at

(a)	$0^{o}C$	(b)	Its critical temperature
(c)	Absolute zero	(d)	Its Boyle temperature

A certain sample of gas has a volume of 0.2 litre measured at 1 atm.

pressure	and	0^{o}	C.	At	the	same	pressure	but	at	273 ⁶	' <i>C</i> ,	its
volume w	vill be						[EAMO	CET 19	992,	93; BH	U 20	005]

(a)	0.4 <i>litres</i>	(b)	0.8 litres
(c)	27.8 <i>litres</i>	(d)	55.6 <i>litres</i>

13.

- 400 cm^3 of oxygen at $27^{\circ}C$ were cooled to $-3^{\circ}C$ without 14 change in pressure. The contraction in volume will be
 - (a) 40 cm^3 (b) 30 cm^3
 - (d) 360 cm^3 (c) 44.4 cm^3
- The pressure p of a gas is plotted against its absolute temperature T15. for two different constant volumes, V_1 and V_2 . When $V_1 > V_2$, the
 - (a) Curves have the same slope and do not intersect
 - (b) Curves must intersect at some point other than T = 0
 - (c) Curve for V_2 has a greater slope than that for V_1

(d) Curve for V_1 has a greater slope than that for V_2

16. Two closed vessels of equal volume containing air at pressure P_1 and temperature T_1 are connected to each other through a narrow tube. If the temperature in one of the vessels is now maintained at T_1 and that in the other at T_2 , what will be the pressure in the vessels

(a)
$$\frac{2P_1T_1}{T_1 + T_2}$$
 (b) $\frac{T_1}{2P_1T_2}$
(c) $\frac{2P_1T_2}{T_1 + T_2}$ (d) $\frac{2P_1}{T_1 + T_2}$

- "One gram molecule of a gas at N.T.P. occupies 22.4 litres." This fact [CPMT 1981, 1995] was derived from
 - (a) Dalton's theory

18.

19.

[CBSE PMT 1989]

- (b) Avogadro's hypothesis
- Berzelius hypothesis (c)
- (d) Law of gaseous volume
- In a closed flask of 5 *litres,* 1.0 g of H_2 is heated from 300 to 600
- K. which statement is not correct [CBSE PMT 1991]
- Pressure of the gas increases (a)
- (b) The rate of collision increases
- The number of moles of gas increases (c)
- The energy of gaseous molecules increases (d)
- Which one of the following statements is false

[Manipal PMT 1991]

[KCET 2005]

[Orissa 1990]

- (a) Avogadro number = 6.02×10^{21}
- (b) The relationship between average velocity (\overline{v}) and root mean square velocity (*u*) is $\overline{v} = 0.9213 u$
- The mean kinetic energy of an ideal gas is independent of the (c) pressure of the gas
- The root mean square velocity of the gas can be calculated by (d) the formula $(3RT/M)^{1/2}$

The compressibility of a gas is less than unity at STP. Therefore [IIT 2000] 20.

- (b) $V_m < 22.4$ litres (a) $V_m > 22.4$ litres
- (d) $V_m = 44.8$ litres (c) $V_m = 22.4$ litres
- In the equation of sate of an ideal gas PV = nRT, the value of the 21. universal gas constant would depend only on
 - The nature of the gas
 - (a) The pressure of the gas (b)
 - The units of the measurement (c)
 - (d) None of these
- In the ideal gas equation, the gas constant R has the dimensions of [NCERT 1982 22. (a) mole-atm K^a (b) *litre mole*
 - litre-atm K mole (d) erg K (c)
- In the equation PV = nRT, which one cannot be the numerical 23. value of R[BIT 1987]
 - (a) $8.31 \times 10^7 erg K^{-1} mol^{-1}$
 - $8.31 \times 10^7 dyne \ cm \ K^{-1} mol^{-1}$ (b)
 - $8.31 JK^{-1}mol^{-1}$ (c)
 - (d) 8.31 *atm*. $K^{-1}mol^{-1}$
- Which one of the following indicates the value of the gas constant R[EAMCET 1 24.
 - (a) 1.987 cal K^a mol^a (b) 8.3 cal K mol (d) 1.987 Joules K mol
 - (c) 0.0821 lit K mol
 - The constant R is
 - (a) Work done per molecule

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25.



	(b) Work done per degree absolute	37.	16 g of oxygen and 3 g of hydrogen are mixed and kept at 760 mm
	(c) Work done per degree per mole(d) Work done per mole		pressure and $0^{o}C$. The total volume occupied by the mixture will be nearly [Vellore CMC 1991]
26.	Select one correct statement. In the gas equation, $PV = nRT$ [CBSE PMT	ľ 1992]	(a) 22.4 <i>litres</i> (b) 33.6 <i>litres</i>
	(a) n is the number of molecules of a gas	-0	(c) 448 litres (d) 44800 ml
	(b) <i>V</i> denotes volume of one mole of the gas	38.	Pure hydrogen sulphide is stored in a tank of 100 <i>litre</i> capacity at
	(c) <i>n</i> moles of the gas have a volume V		$20^{\circ}C$ and 2 <i>atm</i> pressure. The mass of the gas will be [CPMT 1989]
	(d) P is the pressure of the gas when only one mole of gas is		(a) $34 g$ (b) $340 g$ (c) $282 4 g$ (d) $2824 g$
27.	The correct value of the gas constant <i>R</i> is close to	39.	At N.T.P. the volume of a gas is found to be 273 <i>ml</i> . What will be
	[CBSE PMT 1992]		the volume of this gas at 600 mm Hg and $273^{\circ}C$
	(a) 0.082 <i>litre-atmopshere K</i>		[CPMT 1992]
	(b) 0.082 <i>litre-atmosphere</i> K^{-1} mol ⁻¹		(a) 391.8 mL (b) 380 ml
		40	(c) 091.0 m (d) 750 m/ One litre of a gas weighs 2 g at 300 K and 1 atm pressure of the
	(c) 0.082 htte-atmosphere K mole	.	pressure is made 0.75 <i>atm</i> , at which of the following temperatures
	(d) 0.082 $litre^{-1} atmosphere^{-1} K mol$		will one <i>litre</i> of the same gas weigh one gram
28.	S.I. unit of gas constant <i>B</i> is [CPMT 1994]		[CBSE PMT 1992]
20.	(a) 0.0821 litre atm K mole		(a) 430 K (b) 000 K (c) 800 K (d) 900 K
	(b) 2 calories K' mole	41	A wheather balloon filled with hydrogen at 1 atm and $27^{\circ}C$ has
	(c) 8 31 joule K mole	41.	volume equal to 12000 <i>litres.</i> On ascending it reaches a place where
	(d) None		the temperature is $-23^{\circ}C$ and pressure is 0.5 atm. The volume of
~~	(0) for a product of DV = a DT is should be		the balloon is
29.	Gas equation $PV = hKI$ is obeyed by [BHU 2000]		[CBSE PMT 1991]
	(a) Only isothermal process (b) Only adiabatic process (c) $\mathbf{P}_{i}(\mathbf{r}) = 1_{i}(\mathbf{r})$		(a) 24000 <i>litres</i> (b) 20000 <i>litres</i>
20	(c) Both (a) and (b) (d) None of these R		(c) 10000 <i>litres</i> (d) 12000 <i>litres</i>
30.	gas constant R and temperature T is	42.	The density of a gas at $27^{\circ}C$ and 1 <i>atm</i> is <i>d</i> . Pressure remaining
	[AIEEE 2002]		constant at which of the following temperatures will its density
	(a) PT/R (b) PRT		
21	(c) P/RT (d) RT/P		(a) $20^{\circ}C$ (b) $30^{\circ}C$
31.	the pressure must be		(c) 400 K (d) 300 K
	[NCERT 1981; JIPMER 1991]	43.	A sample of gas occupies 100 ml at $27^{\circ}C$ and 740 mm pressure.
	(a) 2 <i>atm</i> (b) 3 <i>atm</i>		When its volume is changed to 80 <i>ml</i> at 740 <i>mm</i> pressure, the temperature of the gas will be
	(c) 4 <i>atm</i> (d) 1 <i>atm</i>		[Vellore CMC 1991]
32.	How many moles of <i>He</i> gas occupy 22.4 <i>litres</i> at $30^{\circ}C$ and one		(a) $21.6^{\circ}C$ (b) $240^{\circ}C$
	atmospheric pressure [KCET 1992]		(a) 21000 (b) 24000 (c)
	$ \begin{array}{c} (a) & 0.90 \\ (b) & 1.0 \\ (c) & 0.11 \\ (d) & 1.0 \\ \end{array} $		(c) $-33^{\circ}C$ (d) $89.5^{\circ}C$
33.	Volume of 0.5 <i>mole</i> of a gas at 1 <i>atm.</i> pressure and $273K$ is	44.	equal to the sum of the partial pressures of the gases under the
00.	[EAMCET 1992]		same conditions is known as [CPMT 1986]
	(a) 22.4 <i>litres</i> (b) 11.2 <i>litres</i>		(a) Boyle's law (b) Charle's law
	(c) 44.8 <i>litres</i> (d) 5.6 <i>litres</i>		(c) Avogadro's law (d) Dalton's law
34.	At $0^{o}C$ and one <i>atm</i> pressure, a gas occupies 100 <i>cc</i> . If the	45.	contain equal number of particles." This statement is a direct
	increased by one-third of absolute temperature, then final volume of		consequence of [Kerala MEE 2002]
	the gas will be		(a) Avogadro's law (b) Charle's law
	[DCE 2000]		(c) Ideal gas equation (d) Law of partial pressure
	(a) 80 cc (b) 88.9 cc	46.	If three unreactive gases having partial pressures P_A , P_B and P_C
35.	Correct gas equation is [CBSE PMT 1989: CPMT 1991]		and their moles are 1, 2 and 3 respectively then their total pressure
00	V_1T_2 V_2T_1 P_1V_1 T_1		D + D + D
	(a) $\frac{1}{P_1} = \frac{2}{P_2}$ (b) $\frac{1}{P_2V_2} = \frac{1}{T_2}$		(a) $P = P_A + P_B + P_C$ (b) $P = \frac{I_A + I_B + I_C}{\epsilon}$
	PT PV VV		6
	(c) $\frac{I_1 I_2}{V} = \frac{I_2 V_2}{T}$ (d) $\frac{V_1 V_2}{TT} = P_1 P_2$		(c) $P = \frac{\sqrt{P_A + P_B + P_C}}{(d) \text{ None}}$
26	v_1 I_2 I_1I_2 Two separate hulbs contain ideal games A and P. The density of $$		3
30.	A is twice that of gas B. The molecular mass of A is half that of gas	47.	Dalton's law of partial pressure will not apply to which of the
	B. The two gases are at the same temperature. The ratio of the		[Bihar MADT 1981]
	pressure of A to that of gas B is		(a) Π_2 and SO_2 (b) Π_2 and Cl_2
	[вни 1994] (а) 2 (b) 1/2		(c) H_2 and CO_2 (d) CO_2 and Cl_2
	(c) 4 (d) $1/4$		

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- 48. Which of the following mixtures of gases does not obey Dalton's law of partial pressure
 - [CBSE PMT 1996: Kerala PMT 2000] (a) O_2 and CO_2 (b) N_2 and O_2
 - (d) NH_3 and HCl(c) Cl_2 and O_2
- To which of the following gaseous mixtures is Dalton's law not 49. applicable

(a)
$$Ne + He + SO_2$$
 (b) $NH_3 + HCl + HBr$

(c) $O_2 + N_2 + CO_2$ (d) $N_2 + H_2 + O_2$

Equal amounts of two gases of molecular weight 4 and 40 are 50. mixed. The pressure of the mixture is 1.1 atm. The partial pressure of the light gas in this mixture is [CBSE PMT 1991]

(a) 0.55 atm (b) 0.11 *atm*

(d) 0.12 atm (c) 1 *atm*

Rate of diffusion of a gas is 51.

52.

- (a) Directly proportional to its density
- (b) Directly proportional to its molecular mass
- (c) Directly proportional to the square root of its molecular mass
- (d) Inversely proportional to the square root of its molecular mass
- Which of the following gas will have highest rate of diffusion

[Pb. CET Sample paper 1993; CPMT 1990]

[IIT 1985; CPMT 1987]

- (a) NH_3 (b) N_2
- (d) O_2 (c) CO_2
- Which of the following relationship is correct, where r is the rate of 53. diffusion of a gas and d is its density [CPMT 1994]

(a)
$$r \propto \sqrt{1/d}$$
 (b) $r \propto \sqrt{d}$

- (c) r = d(d) $r \propto d$
- According to Grahman's law at a given temperature, the ratio of the 54. rates of diffusion r_A / r_B of gases A and B is given by [IIT 1998]
 - (a) $(P_A / P_B)(M_A / M_B)^{1/2}$
 - (b) $(M_A / M_B)(P_A / P_B)^{1/2}$
 - (c) $(P_A / P_B)(M_B / M_A)^{1/2}$
 - (d) $(M_A / M_B)(P_B / P_A)^{1/2}$

(c) 64

(where *P* and *M* are the pressures and molecular weights of gases *A* and *B* respectively)

The ratio of the rate of diffusion of a given element to that of 55. helium is 1.4. The molecular weight of the element is

(a)	2	(b)	4
(c)	8	(d)	16

a∲℃EE^G982] A gas diffuse 1/5 times as fast as hydrogen. Its molecular weight is [CPMT 1992; Bih 56. (a) 50 (b) 25 of X. The molecular weight of X is (c) $25\sqrt{2}$ (d) $50\sqrt{2}$ (a) 64.0

[Kerala PMT 1990]

67.

71.

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The molecular weight of a gas which diffuses through a porous plug 57. at 1/6th of the speed of hydrogen under identical conditions is[EAMCET 1990] (a) 27 (b) 72 70. (c) 36 (d) 48 58. Molecular weight of a gas that diffuses twice as rapidly as the gas

(d) 6.4

- with molecular weight 64 is [EAMCET 1994] (a) 16 (b) 8
- The densities of hydrogen and oxygen are 0.09 and 1.44 $g L^{-1}$. If 59. the rate of diffusion of hydrogen is I then that of oxygen in the same units will be [RPMT 1994]

- If rate of diffusion of A is 5 times that of B, what will be the density 60. ratio of A and B[AFMC 1994]
 - (a) 1/25 (b) 1/5 (d) 4
 - (c) 25
- The densities of two gases are in the ratio of 1 : 16. The ratio of their 61. rates of diffusion is [CPMT 1995] (a) 16:1
 - (b) 4:1 (c) 1:4 (d) 1:16
- 62. At constant volume and temperature conditions, the rate of diffusion D_A and D_B of gases A and B having densities ho_A and ho_B are related by the expression []]T 1993]

(a)
$$D_A = \left[D_B \cdot \frac{\rho_A}{\rho_B} \right]^{1/2}$$
 (b) $D_A = \left[D_B \cdot \frac{\rho_B}{\rho_A} \right]^{1/2}$
(c) $D_A = D_B \left(\frac{\rho_A}{\rho_B} \right)^{1/2}$ (d) $D_A = D_B \left(\frac{\rho_B}{\rho_A} \right)^{1/2}$

Atmolysis is a process of 63.

- (a) Atomising gas molecules
 - (b) The breaking of atoms to sub-atomic particles
 - (c) Separation of gases from their gaseous mixture
- (d) Changing of liquids to their vapour state
- 64. A bottle of ammonia and a bottle of dry hydrogen chloride connected through a long tube are opened simultaneously at both ends, the white ammonium chloride ring first formed will be[11T 1988]
 - (a) At the centre of the tube
 - Near the hydrogen chloride bottle (b)
 - Near the ammonia bottle (c)
 - (d) Throughout the length of the tube
- Which of the following pairs will diffuse at the same rate through a 65. porous plug [EAMCET 1990]

(a)
$$CO, NO_2$$
 (b) NO_2, CO_2

(c)
$$NH_3, PH_3$$
 (d) NOC_2H_6

- 66. If 4 g of oxygen diffuse through a very narrow hole, how much hydrogen would have diffused under identical conditions [CPMT 1971] (a) 16 g (b) 1 g (d) 64 g
 - (c) 1/4 g A gas diffuse at a rate which is twice that of another gas B. The
 - ratio of molecular weights of A to B is EAMCET 1986 (a) 1.0 (b) 0.75
 - (c) 0.50 (d) 0.25
- Two grams of hydrogen diffuse from a container in 10 *minutes*. How 68. many grams of oxygen would diffuse through the same container in the same time under similar conditions [MNR 1980] (a) 0.5 g (b) 4 g
 - (d) 8 g

The rate of diffusion of methane at a given temperature is twice that

- [MNR 1995; Kerala CEE 2001] (b) 32.0
- (d) 80.0 (c) 40.0

X ml of H_2 gas effuses through a hole in a container in 5 seconds. The time taken for the effusion of the same volume of the gas specified below under identical condition is

- []]T 1996]
- (a) 10 seconds : He (b) 20 seconds : O_2

times that of SO_2 at $50^o C$

(c) 25 seconds : CO (d) 55 seconds : CO_2

At what temperature, the rate of effusion of N_2 would be 1.625

[CBSE PMT 1996]

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	(a) 110 <i>K</i> (b) 173 <i>K</i> (c) 373 <i>K</i> (d) 273 <i>K</i>	83.	A closed vessel contains equal number of nitrogen and oxygen molecules at a pressure of <i>P mm</i> . If nitrogen is removed from the
72.	Given the reaction $C(s) + H_2O(l) \rightarrow CO(g) + H_2(g)$ calculate the		system then the pressure will be
	volume of the gases produced at STP from 48.0 g of carbon		[MP PMT 1985]
	(a) 179.2 <i>L</i> (b) 89.6 <i>L</i>		(a) P (b) 2P
	(c) 44.8 <i>L</i> (d) 22.4 <i>L</i>		(c) $P/2$ (d) P^2
73.	4.4 <i>g</i> of a gas at STP occupies a volume of 2.24 <i>L</i> , the gas can be [Hary: (a) O_2 (b) <i>CO</i>	ana (32) ;T	2000]the four tubes of a car are filled to the same pressure with N_2, O_2, H_2 and <i>Ne</i> separately, then which one will be filled first[Manipal PM
	(c) NO_2 (d) CO_2		(a) N_2 (b) O_2
74.	Under what conditions will a pure sample of an ideal gas not only		(c) H_{z} (d) N_{a}
	exhibit a pressure of 1 <i>atm</i> but also a concentration of 1 <i>mole litre</i> -1	85.	Which of the following gas mixture is not applicable for Dalton's law
	$(R = 0.082 litreatmmol^{-1} deg^{-1})$ [CBSE PMT 1003]		or partial pressure $[PB, CEI 2002]$
	(a) At STP		(a) SO_2 and Cl_2 (b) CO_2 and N_2
	(b) When $V = 22.4$ litres		(c) CO and CO_2 (d) CO and N_2
	(b) when $V = 22.4$ miles	86.	At what pressure a quantity of gas will occupy a volume of $ 60 ml$,
	(c) When I = 12 K(d) Impossible under any conditions		if it occupies a volume of 100 <i>ml</i> at a pressure of 720 <i>mm</i> ? (while temperature is constant) : [Pb. CET 2000]
75.	There are 6.02×10^{22} molecules each of N_2, O_2 and H_2 which		(a) 700 mm (b) 800 mm
	are mixed together at 760 mm and 273 K. The mass of the mixture		(c) $100 mm$ (d) $1200 mm$
	in grams is [Pb. PMT 1997]	0-	
	(a) 6.2 (b) 4.12 (c) 3.09 (d) 7	87.	At constant temperature and pressure which gas will diffuse first H_2 or O_2 ? [Pb. CET 2000]
76.	Volume of 4.4 g of CO_2 at NTP is [Pb. CET 1997]		(a) Hydrogen
	(a) 22.4 <i>L</i> (b) 44.8 <i>L</i>		(b) Oxygen
	(c) 2.24 L (d) 4.48 L		(c) Both will diffuse in same time
77.	(a) Decrement (b) Velume	00	(d) None of the above
	(a) Pressure (b) Volume	88.	when a jar containing gaseous mixture or equal volumes or CO_2
78	(c) Number of moles (u) remperature		and H_2 is placed in a solution of sodium hydroxide, the solution
70.	A bottle of cold drift contains 200 <i>m</i> inquite in which CO_2 is 0.		level will [Pb. CET 2001]
	indiar. Suppose CO_2 behaves like all ideal gas, the volume of the		(c) Remain constant (d) Become zero
	dissolved CO_2 at STP is [CBSE PMT 1991]	80	At STP $1g$ CaCO ₂ on decomposition gives CO ₂
	(a) 0.224 litre (b) 0.448 litre	0).	
70	(c) 22.4 <i>ntre</i> (d) 2.24 <i>ntre</i> (d) 2.24 <i>ntre</i>		(a) 22.4 litre (b) 2.24 litre
79.	this gas at N.T.P. is		(c) 0.224 litre (d) 11.2 litre
	[MNR 1982; CBSE PMT 1991]	90.	At NTP, the density of a gas, whose molecular weight is 45 is
	(a) 1 <i>L</i> (b) 11.2 <i>L</i>		[Pb. CET 2001, 03]
_	(c) 22.4 <i>L</i> (d) 20 <i>L</i>		(a) 44.8 gm/litre (b) 11.4 gm/litre
80.	A pre-weighed vessel was filled with oxygen at N.T.P. and weighted.		(c) 2 gm/litre (d) 3 gm/litre
	It was then evacuated, filled with SO_2 at the same temperature	91. 1	what is the ratio of diffusion rate of oxygen and hydrogen
	and pressure, and again weighted. The weight of oxygen will be NCERT	[1989]	(a) 1:4 (b) 4:1
	(a) The same as that of SO_2		(c) 1:8 (d) 8:1
	(b) $\frac{1}{2}$ that of SQ ₂	92.	The maximum number of molecules is present in
	$\frac{1}{2}$		[CBSE PMT 2004]
	(c) Twice that of SO_2		(a) 0.5 g of H_2 gas (b) 10 g of O_2 gas (c) 15 L of H_2 gas at STP (d) 5 L of N_2 gas at STP
	(d) One fourth that of SO_2	93.	One litre oxygen gas at STP will weigh [Ph. CET 2004]
81.	Five grams each of the following gases at $87^{\circ}C$ and 750 mm	-00	(a) 1.43 g (b) 2.24 g
	pressure are taken. Which of them will have the least volume[MNR 199	1]	(c) 11.2 g (d) 22.4 g
	(a) <i>HF</i> (b) <i>HCl</i>	94.	How will you separate mixture of two gases [AFMC 2004]
	(c) <i>HBr</i> (d) <i>HI</i>		(a) Fractional distillation technique (b) Crahama law of diffusion technicus
82.	Who among the following scientists has not done any important		(c) Osmosis
	worк on gases [Bihar MADT 1980]		(d) Chromatography
	(a) boyie (b) Charles (c) Avogadro (d) Faraday	95.	The rate of diffusion of hydrogen gas is
	(c) Avogaulo (u) Faladay		[MH CET 2003; Pb. CET 2000]
			(a) 1.4 times to He gas (b) Same as He gas

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	(c) 5 times to He gas (d)) 2 times to <i>He</i> gas
96.	Hydrogen diffuses six times faster the gas A is	nan gas A . The molar mass of [KCET 2004]
	(a) 72 (b)) 6
	(c) 24 (d)) 36
9 7.	At what pressure will a quantity of	gas, which occupies $100 ml$ at
	a pressue of $720mm$, occupy a vol	ume of 84 <i>ml</i> [DPMT 2004]
	(a) 736.18 mm (b)) 820.20 <i>mm</i>
08	(c) $784.15 mm$ (d)) 857.14 <i>mm</i>
90.	temperature of A are all twice	that of $B_{\rm c}$ then the ratio of
	number of molecules of A and B	are [AFMC 2004]
	(a) 1:2 (b)) 2
	(c) 1:4 (d)) 4
99.	A mixture of NO_2 and N_2O_4 h	as a vapour density of 38.3 at
	300K. What is the number of me	oles of NO_2 in $100g$ of the
	mixture $(x) = 0.042$ (b)	[Kerala PMT 2004]
	(c) 3.4 (d)) 3.86
	(e) 0.437	
100.	A cylinder of 5 litres capacity, filled with another evacuated cylinder	d with air at NTP is connected
	resultant air pressure in both the cyl	inders will be [BHU 2004]
	(a) 10.8 cm of Hg (b)) 14.9 <i>cm</i> of <i>Hg</i>
101.	A certain mass of gas occupies a vol	ume of 300 c.c. at 27 <i>C</i> and 620
	<i>mm</i> pressure. The volume of this	gas at $47^{\circ}C$ and 640 mm
	pressure will be	[MH CET 2003]
	(a) 400 c.c. (b) $(c) 310 c.c.$ (d)) 510 c.c.
102.	What will be the volume of the mixtu	are after the reaction?
	$NH_{3} + HCl \rightarrow NH_{4}Cl$ $4 \text{ litre} \xrightarrow{1.5 \text{ litre}} (solid)$	[BVP 2004]
	(a) 0.5 litre (b)) 1 litre
	(c) 2.5 litre (d)) 0.1 litre
103.	The pressure and temperature of 4 doubled. Then the volume of carbon	<i>dm</i> [°] of carbon dioxide gas are dioxide gas would be[KCET 2004]
	(a) $2 dm^3$ (b)) $3dm^3$
	(c) $4 dm^3$ (d)) $8 dm^3$
104.	If the absolute temperature of an	ideal gas become double and
	pressure become half, the volume of	gas would be
	(a) Bemain unchange (b)	[Kerala CET 2005]
	(c) Will be four time (d)) will be half
	(e) Will be one fourth	
105.	At what temperature, the sample o double of its pressure, if the initial	f neon gas would be heated to volume of gas is/are reduced to
	15% at $75^{o}C$	[Kerala CET 2005]
	(a) $319^{\circ}C$ (b)) $592^{\circ}C$
	(c) $128^{\circ}C$ (d)) $60^{\circ} C$
	(e) $90^{\circ}C$	
106.	Equation of Boyle's law is	[DPMT 2005]
	dP dV	dP = dV
	(a) $= =$ (b)) = +
	(a) $\frac{1}{p} = -\frac{1}{V}$ (b)	$\frac{1}{P} = +\frac{1}{V}$
	(a) $\frac{1}{p} = -\frac{1}{V}$ (b)	$\frac{1}{P} = +\frac{1}{V}$ $\frac{d^2 P}{d^2 V} = +\frac{d^2 V}{d^2 V}$

Kinetic molecular theory of gases and Molecular collisions

	Postulate of kinetic theory is	[EAMCET 1980]	
	(a) Atom is indivisible		
	(b) Gases combine in a simp	le ratio	
	(c) There is no influence of	gravity on the molecules of a gas	
	(d) None of the above		
2.	According to kinetic theory of	gases, [EAMCET 1980]	
	(a) There are intermolecular	attractions	
	(b) Molecules have considera	ble volume	
	(c) No intermolecular attrac	ions	
	(d) The velocity of molecules	decreases after each collision	
3.	In deriving the kinetic gas ec square velocity of the molecul	uation, use is made of the root mean es because it is	
		[Bihar MADT 1980]	
	(a) The average velocity of t	ne molecules	
	(b) The most probable veloc	ty of the molecules	
	(c) The square root of the a	verage square velocity of the molecules	
	(d) The most accurate form calculations	in which velocity can be used in these	
4.	Kinetic energy of a gas depend	ls upon its[Bihar MADT 1982]	
	(a) Molecular mass	(b) Atomic mass	
	(c) Equivalent mass	(d) None of these	
5.	The kinetic theory of gases	perdicts that total kinetic energy of a	
	gaseous assembly depends on	[NCERT 1984]	
	(a) Pressure of the gas		
	(b) Temperature of the gas		
	(c) Volume of the gas		
~	(d) Pressure, volume and ter	perature of the gas	
6.	According to kinetic theory of	gases, the energy per mole of a gas is	
	(a) 1.5 BT		
	(-)		
	(c) 0.5 <i>RT</i>	(d) $2.5 RT$	
7.	(c) 0.5 <i>RT</i> Internal energy and pressure of	(d) 2.5 <i>RT</i> (d) 2.5 <i>RT</i> of a gas per unit volume are related as[CBSE PA	ИТ
7.	(c) 0.5 <i>RT</i> Internal energy and pressure (c) $P = \frac{2}{2}F$	(b) RT (d) 2.5 RT of a gas per unit volume are related as [CBSE PA (b) $P = \frac{3}{E}$	ИТ
7.	(c) 0.5 <i>RT</i> Internal energy and pressure of (a) $P = \frac{2}{3}E$	(b) RT (d) 2.5 RT of a gas per unit volume are related as [CBSE PA (b) $P = \frac{3}{2}E$	ИТ
7.	(c) 0.5 <i>RT</i> Internal energy and pressure of (a) $P = \frac{2}{3}E$	(b) RT (d) 2.5 RT of a gas per unit volume are related as [CBSE PA (b) $P = \frac{3}{2}E$	νт
7.	(c) 0.5 <i>RT</i> Internal energy and pressure (a) $P = \frac{2}{3}E$ (c) $P = \frac{1}{2}E$	(b) P (c) $P = \frac{3}{2}E$ (c) $P = 2E$	ИТ
7.	(c) 0.5 <i>RT</i> Internal energy and pressure of (a) $P = \frac{2}{3}E$ (c) $P = \frac{1}{2}E$ The translational kinetic energy	(b) RT (d) 2.5 RT of a gas per unit volume are related as [CBSE PA (b) $P = \frac{3}{2}E$ (d) $P = 2E$ v of an ideal gas depends only on its	ИТ
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12. According to kinetic theory of gases, for a diatomic molecule

[MNR 1991]

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- (a) The pressure exerted by the gas is proportional to the mean velocity of the molecules
- The pressure exerted by the gas is proportional to the root (b) mean square velocity of the molecules
- The root mean square velocity is inversely proportional to the (c) temperature
- The mean translational kinetic energy of the molecules is (d) proportional to the absolute temperature
- 13. At STP, 0.50 mol H_2 gas and 1.0 mol He gas

[CBSE PMT 1993, 2000]

- (a) Have equal average kinetic energies
- (b) Have equal molecular speeds
- Occupy equal volumes (c)
- Have equal effusion rates (d)
- Which of the following expressions correctly represents the 14. relationship between the average molar kinetic energy, K.E., of CO and N_2 molecules at the same temperature

[CBSE PMT 2000]

- $\overline{KE}_{CO} = \overline{KE}_{N_2}$ (a)
- (b) $\overline{KE}_{CO} > \overline{KE}_{N_2}$
- (c) $\overline{KE}_{CO} < \overline{KE}_{N_2}$

15.

- (d) Cannot be predicted unless the volumes of the gases are given
- Indicate the correct statement for a 1-L sample of $N_2(g)$ and $CO_2(g)$ at 298 K and 1 atm pressure
 - (a) The average translational KE per molecule is the same in N_2 and CO_2
 - (b) The *rms* speed remains constant for both N_2 and CO_2
 - (c) The density of N_2 is less than that of CO_2
- The total translational KE of both N_2 and CO_2 is the same
- 16. With increase of pressure, the mean free path

[Pb. CET 1985]

- (a) Decreases (b) Increases (c) Does not change (d) Becomes zero
- Which one of the following statements is NOT true about the effect 17. of an increase in temperature on the distribution of molecular speeds in a gas AIEEE 2005
 - (a) The most probable speed increases
 - (b) The fraction of the molecules with the most probable speed increases
 - (c) The distribution becomes broader
 - (d) The area under the distribution curve remains the same as under the lower temperature
- 18. If P, V, M, T and R are pressure, volume, molar mass, temperature and gas constant respectively, then for an ideal gas, the density is given by [CBSE PMT 1989, 91]

a)
$$\frac{RT}{PM}$$
 (b) $\frac{P}{RT}$

c)
$$\overline{V}$$
 (d) \overline{RT}

- An ideal gas will have maximum density when [CPMT 2000] 19. (a) P = 0.5 atm, T = 600 K
 - P = 2 atm, T = 150 K(h)
 - P = 1 atm, T = 300 K(c)
 - P = 1.0 atm, T = 500 K(d)
- If the inversion temperature of a gas is $-80^{o}C$, then it will 20. produce cooling under Joule-Thomson effect at

- (a) 298 K (b) 273 K (c) 193 K (d) 173 K
- Ratio of C_p and C_y of a gas 'X is 1.4. The number of atoms of the 21. gas 'X present in 11.2 *litres* of it at N.T.P. is
- [CBSE PMT 1989] (b) 1.2×10^{24} (a) 6.02×10^{23} (c) 3.01×10^{23} (d) 2.01×10^{23} The density of air is 0.00130 g/ml. The vapour density of air will be[DCE 2000] 22. (a) 0.00065 (b) 0.65 (c) 14.4816 (d) 14.56 At $100^{\circ}C$ and 1 *atm*, if the density of liquid water is 1.0 g cm⁻³ 23. and that of water vapour is 0.0006 $g\ m^{-3}$, then the volume occupied by water molecules in 1 litre of steam at that temperature [IIT 2000] is (b) 60 cm³ (a) $6 \ cm^3$ (c) 0.6 cm^3 (d) 0.06 cm^3 The ratio γ for inert gases is [AFMC 1990] 24. (a) 1.33 (b) 1.66 (d) 1.99 (c) 2.13 The density of neon will be highest at 25. [CBSE PMT 1990] (a) S.T.P. (b) $0^{\circ}C.2 atm$ (c) $273^{\circ}C, 1 atm$ (d) $273^{\circ}C, 2 atm$ 26. Absolute zero is defined as the temperature [CBSE PMT 1990] (a) At which all molecular motion ceases (b) At which liquid helium boils (c) At which ether boils (d) All of the above 27. Consider the following statements : (1) Joule-Thomson experiment is isoenthalpic as well as adiabatic. (2)A negative value of μ_{JT} (Joule Thomson coefficient corresponds to warming of a gas on expansion. The temperature at which neither cooling nor heating effect is (3)
 - observed is known as inversion temperature. Which of the above statements are correct
 - (a) 1 and 2 (b) 1 and 3
 - (c) 2 and 3 (d) 1, 2 and 3
 - Vibrational energy is [Pb. CET 1985]
 - (a) Partially potential and partially kinetic
 - (b) Only potential
 - Only kinetic (c)

28.

- (d) None of the above
- At the same temperature and pressure, which of the following gases 29. will have the highest kinetic energy per mole

(b) Oxygen

[MNR 1991]

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(a) Hydrogen (c) Methane (d) All the same

30. Dimensions of pressure are the same as that of

				[CB3E PMT 1995]
	(a)	Energy	(b)	Force
	(c)	Energy per unit volume	(d)	Force per unit volume
31.	The	density of a gas An is	three ti	imes that of a gas <i>B</i> . if the
	mol	ecular mass of A is M, the	molecula	ar mass of <i>B</i> is
				[CPMT 1987]
	(a)	3 <i>M</i>	(b)	$\sqrt{3}$ M
	(c)	<i>M</i> / 3	(d)	$M/\sqrt{3}$

Molecular speeds



- The ratio of root mean square velocity to average velocity of gas 1. molecules at a particular temperature is [IIT 1981] (a) 1.086 : 1 (b) 1:1.086
 - (c) 2:1.086 (d) 1.086 : 2

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- Which is not true in case of an ideal gas
- (a) It cannot be converted into a liquid
 - (b) There is no interaction between the molecules
 - (c) All molecules of the gas move with same speed
 - At a given temperature, PV is proportional to the amount of (d) the gas

[CBSE PMT 1991]

The ratio among most probable velocity, mean velocity and root 3 mean square velocity is given by [CBSE PMT 1993]

(a)
$$1:2:3$$
 (b) $1:\sqrt{2}:\sqrt{3}$

- (d) $\sqrt{2}: \sqrt{8/\pi}: \sqrt{3}$ (c) $\sqrt{2}: \sqrt{3}: \sqrt{8/\pi}$
- Which of the following has maximum root mean square velocity at 4 [Manipal PMT 2002] the same temperature
 - (b) *CO*₂ (a) SO_2 (c) O_2 (d) H_2
- The temperature at which RMS velocity of SO_2 molecules is half 5 that of He molecules at 300 K is [NTSE 1991] (h) 600 V(a) 150 K

(a)	150 N	(0)	000 /
(c)	900 K	(d)	1200 K

- At $27^{\circ}C$, the ratio of *rms* velocities of ozone to oxygen is 6. [EAMCET 1992]
 - (a) $\sqrt{3/5}$ (b) $\sqrt{4/3}$ (c) $\sqrt{2/3}$
 - (d) 0.25
- The average kinetic energy of an ideal gas per molecule in SI units at 7. $25^{o}C$ will be [CBSE PMT 1996]
 - (a) $6.17 \times 10^{-21} kJ$ (b) $6.17 \times 10^{-21} J$ (c) $6.17 \times 10^{-20} J$ (d) $7.16 \times 10^{-20} J$
 - At what temperature the RMS velocity of SO_2 be same as that of
 - O₂ at 303 K (a) 273 K (b) 606 K
 - (c) 303 K (d) 403 K Among the following gases which one has the lowest root mean
 - square velocity at $25^{\circ}C$ [EAMCET 1983]
 - (a) SO_2 (b) N_2
 - (d) *Cl*₂ (c) O_2
- The root mean square velocity of an ideal gas in a closed container 10. of fixed volume is increased from $5 \times 10^4 \ cm \ s^{-1}$ to

 $10 \times 10^4 \ cm \ s^{-1}$. Which of the following statement correctly explains how the change is accomplished

[Pb. CET 1986]

[KCET 2001]

- (a) By heating the gas, the temperature is doubled
- (b) By heating the gas, the pressure is quadrupled (*i.e.* made four times)
- (c) By heating the gas, the temperature is quadrupled
- (d) By heating the gas, the pressure is doubled
- The rms velocity at NTP of the species can be calculated from the 11. [EAMCET 1990] expression

(a)
$$\sqrt{\frac{3P}{d}}$$
 (b) $\sqrt{\frac{3PV}{M}}$

(c)
$$\sqrt{\frac{3 RT}{M}}$$
 (d) All the above

12. Root mean square velocity of a gas molecule is proportional to [CBSE PMT 1990]

- (a) $m^{1/2}$ (b) m^0
- (c) $m^{-1/2}$ (d) *m*
- At constant volume, for a fixed number of moles of a gas, the 13. pressure of the gas increases with increase in temperature due to[11T 1992] (a) Increase in the average molecular speed
 - Increased rate of collision amongst molecules (b)
 - (c) Increase in molecular attraction
 - (d) Decrease in mean free path
- Molecular velocities of the two gases at the same temperature are 14. u_1 and u_2 . Their masses are m_1 and m_2 respectively. Which of the following expressions is correct

[BHU 1994]

(a)
$$\frac{m_1}{u_1^2} = \frac{m_2}{u_2^2}$$
 (b) $m_1 u_1 = m_2 u_2$
(c) $\frac{m_1}{u_1} = \frac{m_2}{u_2}$ (d) $m_1 u_1^2 = m_2 u_2^2$

- The temperature of the gas is raised from $27^{o}C$ to $927^{o}C$, the 15. root mean square velocity is [CBSE PMT 1994]
 - (a) $\sqrt{927/27}$ times the earlier value
 - (b) Same as before
 - (c) Halved
 - (d) Doubled
- The ratio between the root mean square velocity of H at 50 K and 16. that of O_2 at 800 K is [IIT 1996]

The root mean square velocity of an ideal gas at constant pressure 17. varies density (d) as [IIT 2000]

(a)
$$d^2$$
 (b) d
(c) \sqrt{d} (d) $1/\sqrt{d}$

- Consider a mixture of SO_2 and O_2 kept at room temperature. 18. Compared to the oxygen molecule, the SO_2 molecule will hit the wall with (a) Smaller average speed (b) Greater average speed
 - (d) Greater mass (c) Greater kinetic energy
- The *rms* speed of N_2 molecules in a gas is *u*. If the temperature is 19. doubled and the nitrogen molecules dissociate into nitrogen atoms, the *rms* speed becomes

(a)
$$u/2$$
 (b) $2u$
(c) $4u$ (d) $14u$

- Choose the correct arrangement, where the symbols have their usual 20. meanings
 - (a) $\overline{u} > u_p > u_{rms}$ (b) $u_{rms} > \overline{u} > u_p$ (c) $u_p > \overline{u} > u_{rms}$ (d) $u_p > u_{rms} > \overline{u}$
- The ratio of most probable velocity to that of average velocity is [JEE Orissa 200 21. (a) $\pi/2$ (b) $2/\pi$
 - (c) $\sqrt{\pi}/2$ (d) $2/\sqrt{\pi}$
- The r.m.s. velocity of a certain gas is v at 300K. The 22. temperature, at which the r.m.s. velocity becomes double [Pb. CET 2002]
 - (a) 1200*K* (b) 900 K
 - (c) 600*K* (d) 150K
- The r.m.s. velocity of a gas depends upon [DCE 2002] 23.
 - (a) Temperature only

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- (b) Molecular mass only
- Temperature and molecular mass of gas (c)
- (d) None of these
- What is the pressure of 2 mole of NH_3 at $27^{\circ}C$ when its 24. volume is 5 litre in vander Waal's equation (a = 4.17, b = 0.03711)[JEE Orissa-2004] (a) 10.33 atm (b) 9.33 atm (c) 9.74 atm (d) 9.2 atm
- 25. The root mean square velocity of one mole of a monoatomic having molar mass M is U_{ms} . The relation between the average kinetic energy (E) of the U_{mus} is

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16.

(a)
$$U_{ms} = \sqrt{\frac{3E}{2M}}$$
 (b) $U_{ms} = \sqrt{\frac{2E}{3M}}$
(c) $U_{ms} = \sqrt{\frac{2E}{M}}$ (d) $U_{ms} = \sqrt{\frac{E}{3M}}$

Ratio of average to most probable velocity is 26.

		0	•		5	[Orissa JEE 2005]
(a)	1.128			(b)	1.224	
(c)	1.0			(d)	1.112	

If the v_{ms} is $30R^{1/2}$ at 27^oC then calculate the molar mass of 27. gas in kilogram. [DPMT 2005] (a) 1 (b) 2 (c) 4 (d) 0.001

Real gases and Vander waal's equation

1.	The Vander Waal's equation	on explains the behaviour of	
			[DPMT 1981]
	(a) Ideal gases	(b) Real gases	

(a)	lucal gases	(\mathbf{U})	fical gases
(c)	Vapour	(d)	Non-real gases

Gases deviate from the ideal gas behaviour because their molecules [NCERT 1981] 2. Possess negligible volume (a)

- (b) Have forces of attraction between them
- (c) Are polyatomic
- (d) Are not attracted to one another
- The compressibility factor of a gas is defined as Z = PV/RT. The з. compressibility factor of ideal gas is ----

In Vander Waal's equation of state for a non-ideal gas, the term that 4. accounts for intermolecular forces is [CBSE PMT 1990; IIT 1988]

(a)
$$(V-b)$$
 (b) $(RT)^{-1}$
(c) $\left(P + \frac{a}{V^2}\right)$ (d) RT

Vander Waal's equation of state is obeyed by real gases. For *n* moles 5. of a real gas, the expression will be

[IIT 1992; Pb. CET 1986; DPMT 1986]

(a)
$$\left(\frac{P}{n} + \frac{na}{V^2}\right) \left(\frac{V}{n-b}\right) = RT$$

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

6. Any gas shows maximum deviation from ideal gas at

(a) $0^{o} C$ and 1 atmospheric pressure

- (b) $100^{\circ}C$ and 2 atmospheric pressure
- $-100^{\circ}C$ and 5 atmospheric pressure (c)
- $500^{\circ}C$ and 1 atmospheric pressure (d)

The temperature at which the second virial coefficient of real gas is zero is called [AFMC 1993] (a) Critical temperature

- (b) Eutetic point
- (c)Boiling point
- (d) Boyle's temperature

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When is deviation more in the behaviour of a gas from the ideal gas
equation PV = nRT
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[DPMT 1981; NCERT 1982; CBSE PMT 1993]

[CPMT 1996]

- (a) At high temperature and low pressure
- (b) At low temperature and high pressure
- (c) At high temperature and high pressure
- (d) At low temperature and low high pressure
- Vander Waal's constants 'a' and 'b' are related with respectively [RPMT 1994]
 - (a) Attractive force and bond energy of molecules
 - (b) Volume and repulsive force of molecules
 - Shape and repulsive forces of molecules (c)
 - Attractive force and volume of the molecules (d)
- Gas deviates from ideal gas nature because molecules

 - (a) Are colourless
 - (b) Attract each other
 - Contain covalent bond (c)
 - (d) Show Brownian movement
 - The Vander Waal's equation reduces itself to the ideal gas equation
- [Kerala MEE 2001; CBSE PMT 2002] at (a) High pressure and low temperature
- (b) Low pressure and low temperature
- Low pressure and high temperature (c)
- (d) High pressure and high temperature
- The compressibility factor for an ideal gas is
- [IIT 1997] (a) 1.5 (b) 1.0
 - (c) 2.0 (d) ∞
- 13. When an ideal gas undergoes unrestrained expansion, no cooling occurs because the molecules [IIT 1984, 89]
 - (a) Are above the inversion temperature
 - (b) Exert no attractive force on each other
 - (c) Do work equal to loss in kinetic energy
 - (d) Collide without loss of energy
 - A gas is said to behave like an ideal gas when the relation PV/T = constant. When do you expect a real gas to behave like an ideal gas

[IIT 1999; CBSE PMT 1990; CPMT 1991]

- (a) When the temperature is low
- (b) When both the temperature and pressure are low
- When both the temperature and pressure are high (c)
- (d) When the temperature is high and pressure is low
- A real gas most closely approaches the behaviour of an ideal gas at[KCET 1992] 15.
 - (a) 15 *atm* and 200 *K* (b) 1 *atm* and 273 *K*
 - (d) 15 atm and 500 K (c) 0.5 *atm* and 500 *K*
 - The temperature at which real gases obey the ideal gas laws over a
 - wide range of pressure is called
 - (a) Critical temperature
 - (b) Boyle temperature
 - (c) Inversion temperature

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[AFMC 1993; IIT 1981, 94]

[CPMT 1991]

(d) Reduced temperature

17. At low pressure, the Vander Waal's equation is reduced to

(a)
$$Z = \frac{pV_m}{RT} = 1 - \frac{ap}{RT}$$
 (b) $Z = \frac{pV_m}{RT} = 1 + \frac{b}{RT}p$
(c) $pV_m = RT$ (d) $Z = \frac{pV_m}{RT} = 1 - \frac{a}{RT}$

18. At high temperature and low pressure, the Vander Waal's equation is reduced to

(a)
$$\left(p + \frac{a}{V_m^2}\right)(V_m) = RT$$

(b)
$$pV_m = RT$$

(c)
$$p(V_m - b) = RT$$

(d)
$$\left(p + \frac{a}{V_m^2}\right)(V_m - b) = RT$$

- When helium is allowed to expand into vacuum, heating effect is observed. Its reason is that [CPMT 1987]
 - $(a) \quad \text{Helium is an ideal gas} \\$
 - (b) Helium is an inert gas
 - (c) The inversion temperature of helium is very low
 - $\left(d\right)$ $% \left(d\right)$ The boiling point of helium is the lowest among the elements
- 20.
 In van der Waal's equation of state of the gas law, the constant 'b' is a measure of

 [AIEEE 2004]
 - (a) Volume occupied by the molecules
 - (b) Intermolecular attraction
 - (c) Intermolecular repulsions
 - (d) Intermolecular collisions per unit volume
- **21.** In which molecule the vander Waal's force is likely to be the most important in determining the *m.pt.* and *b.pt.*

[DPMT 2000]

[BHU 1984]

[CBSE PMT 1992]

(a)	H_2S	(b)	Br_2

- (c) HCl (d) CO
- **22.** Pressure exerted by 1 mole of methane in a 0.25 litre container at 300K using vander Waal's equation (given $1 = 2.253 atml^2 mol^{-2}, b = 0.0428 litmol^{-1}$) is

[Orissa JEE 2005]

 (a)
 82.82 atm
 (b)
 152.51 atm

 (c)
 190.52 atm
 (d)
 70.52 atm

Critical state and Liquefaction of gases

- Which set of conditions represents easiest way to liquefy a gas[NCERT 1983]
 (a) Low temperature and high pressure
 (b) 11 11 11
 - (b) High temperature and low pressure
 - (c) Low temperature and low pressure
 - (d) High temperature and high pressure
 - Adiabatic demagnetisation is a technique used for
 - (a) Adiabatic expansion of a gas
 - (b) Production of low temperature
 - (c) Production of high temperature
 - (d) None

2.

3. An ideal gas can't be liquefied because

(a) Its critical temperature is always above $0^{o}C$

- (b) Its molecules are relatively smaller in size
- (c) It solidifies before becoming a liquid
- (d) Forces operative between its molecules are negligible
- However great the pressure, a gas cannot be liquefied above its
 - (a) Boyle temperature

4

- (b) Inversion temperature
- (c) Critical temperature
- (d) Room temperature
- 5. An ideal gas obeying kinetic theory of gases can be liquefied if[CBSE PMT 1995]
 - (a) Its temperature is more than critical temperature T_c
 - (b) Its pressure is more than critical pressure P_c
 - (c) Its pressure is more than P_c at a temperature less than T_c
 - (d) It cannot be liquefied at any value of P and T

6. The Vander Waal's parameters for gases W, X, Y and Z are b (L mol Gas a (atm L' mol·) w 4.0 0.027 х 8.0 0.030 γ 6.0 0.032 Z 12.0 0.027 Which one of these gases has the highest critical temperature (a) W (b) X (d) (c) *Y* ZThe Vander Waal's constant 'a' for the gases O_2, N_2, NH_3 and 7. CH_4 are 1.3, 1.390, 4.170 and 2.253 $L^2 atmmol^{-2}$ respectively. The gas which can be most easily liquefied is [IIT 1989] (a) O_2 (b) N_{2} (c) NH_3 (d) CH_A 8. A gas can be liquefied [AFMC 2005] (a) Above its critical temperature At its critical temperature (b) (c) Below its critical temperature (d) At any temperature 9. Which of the following is correct for critical temperature (a) It is the highest temperature at which liquid and vapour can coexist Beyond the critical temperature, there is no distinction between (b) the two phases and a gas cannot be liquefied by compression (c) At critical temperature (T_c) the surface tension of the system is zero (d) At critical temperature the gas and the liquid phases have different critical densities A gas has a density of 2.68 g/L at stp. Identify the gas 10. (a) NO_2 (b) *Kr* (c) COS (d) SO_2 Weight of 112 *ml* of oxygen at NTP on liquefaction would be[DPMT 1984] (a) 0.32 g (b) 0.64 g (c) 0.16 g (d) 0.96 g Critical Thinking **Objective Questions** As the temperature is raised from $20^{\circ}C$ to $40^{\circ}C$ the average 1.

kinetic energy of neon atoms changes by a factor of which of the following [AIEEE 2004]

(a) 313/293

CLICK HERE

(b) $\sqrt{(313/293)}$

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	(c) 1/2	(d) 2	11.	If C_1, C_2, C_3 represent	the speeds of n_1, n_2, n_3 molec	ules,
2.	A gas is found to have a for	mula $[CO]_x$. If its vapour density is 70,		then the root mean square sp	eed is [IIT 1	993]
	the value of x is	[DCE 2004]		$\left(n_{1}C^{2}+n_{2}C^{2}+n_{3}C^{2}\right)$	$(1/2)^{1/2}$	
	(a) 2.5	(b) 3.0 (d) 6.0		(a) $\left \frac{n_1 c_1 + n_2 c_2 + n_3 c_3}{n_1 + n_2 + n_2 + n_3 c_4} \right $	<u> </u>	
3.	Which of the given sets of te	mperature and pressure will cause a gas		$(1 C^2 + m C^2 + m C^2)$	1/2	
	(a) 100° C and 4 atm	(b) 100° C and 2 atm		(b) $\frac{(n_1C_1 + n_2C_2 + n_3C_3)}{n_1 + n_2 + n_3 + \dots}$	+) 	
	(c) $-100^{\circ} C$ and 4 atm	(d) $0^{\circ}C$ and 2 atm		$(n_1C_1^2)^{1/2}$ $(n_2C_2^2)^{1/2}$	$(n_3C_3^2)^{1/2}$	
4.	The molecular weight of O_2	and SO_2 are 32 and 64 respectively.		(c) $\frac{n_1}{n_1} + \frac{n_2}{n_2}$	$-+\frac{1}{n_3}+\frac{1}{n_3}$	
	If one <i>litre</i> of O_2 at 15^o	C and 750 mm pressure contains 'N		$\int (n_1C_1 + n_2C_2 + n_2C_2)$	$+)^{2}]^{1/2}$	
	molecules, the number of mo	blecules in two <i>litres</i> of SO_2 under the		(d) $\frac{(1 + n_2)^2}{(n_1 + n_2) + n_3 +}$)	
	same conditions of temperate	are and pressure will be [CBSE 1990; MNR 1	991] ₁₂	E0 m/ of hydrogen diffuses or	' I through a small hale from a vess	al in
	(a) N / 2	(b) <i>N</i>	12.	20 <i>minutes</i> . The time needed	for 40 <i>ml</i> of oxygen to diffuse out	is[CBSE PMT 19
	(c) 2 <i>N</i>	(d) 4 <i>N</i>		(a) 12 <i>min</i>	(b) 64 <i>min</i>	
5.	What is the relationship betw	veen the average velocity (v), root mean		(c) 8 <i>min</i>	(d) 32 min	
	square velocity (u) and most	probable velocity (a)	13.	At what temperature will the	average speed of CH_4 molecules	have
		[AFMC 1994]		the same value as O_2 has at	300 K	
	(a) $\alpha: v: u:: 1: 1.128: 1$.224			[CBSE PMT 1	989]
	(b) $\alpha: v: u:: 1.128: 1: 1$.224		(a) 1200 <i>K</i>	(b) 150 <i>K</i>	
	(c) $\alpha: v: u:: 1.128: 1.22$	24 : 1		(c) $600 K$	(d) 300 <i>K</i>	
	(d) $\alpha : y : y : 1 = 124 \cdot 127$	28 • 1	14.	A sample of O_2 gas is c	ollected over water at $23^{\circ}C$	at a
6	(u) $u \cdot v \cdot u \dots 1 \cdot 124 \cdot 124$	ments · For diatomic gases the ratio		barometric pressure of 751 a	mm Hg (vapour pressure of wate	r at
0.	C_p / C_v is equal to			$23^{o}C$ is 21 <i>mm Hg</i>). The sample collected is	partial pressure of O_2 gas in [CBSE PMT 1	the 993]
	(1) 1.40 (lower temperature)			(a) 21 <i>mm Hg</i>	(b) 751 <i>mm Hg</i>	-
	(2) 1.66 (moderate temperate	ıre)		(c) 0.96 <i>atm</i>	(d) 1.02 <i>atm</i>	
	(3) 1.29 (higher temperature)		15.	In an experiment during the	analysis of a carbon compound, 14	5 l
	which of the above statemen	ts are correct		of H_2 was collected at	760 mm Hg pressure and 27	^o C
	(a) $I, 2 and 3$	(b) $1 \text{ and } 2$		temperature. The mass of H_{c}	is nearly	
7	The compressibility factor for	(u) Tanu S ran ideal gas is [MP PFT 2004]			ÍMNR 1	987]
<i>/</i> ·	(a) 1.5	(b) 1.0		(a) 10 g	(b) 12 g	5-11
	(c) 2.0	(d) ∞		(c) $24 g$	(d) 6 g	
8.	The compressibility factor o	f a gas is less than 1 at STP. Its molar	16.	The volume of $1 g$ each of	methane (CH_A) , ethane (C_2H)	I_6),
	volume V_m will be	[MP PET 2004]		propane (C_2H_8) and butane	(C_4H_{10}) was measured at 350	о <i>к</i>
	(a) $V_m > 22.42$	(b) $V_m < 22.42$		and 1 <i>atm</i> . What is the volu	ame of butane [NCERT	1981]
	(c) $V_m = 22.42$	(d) None		(a) 495 cm^3	(b) 600 <i>cm</i> ³	
9.	If some <i>moles</i> of O_2 diffuse	e in 18 <i>sec</i> and same <i>moles</i> of other gas		(c) 900 cm^3	(d) 1700 cm^3	
	diffuse in 45 <i>sec</i> then what gas	is the molecular weight of the unknown [CPMT 1988]	17.	The ratio of the rate of dif	fusion of helium and methane ur and temperature will be [11] 2	nder 0051
	45 ²	18^{2}		(a) 4	(b) 2	
	(a) $\frac{43}{18^2} \times 32$	(b) $\frac{10}{45^2} \times 32$		(c) 1	(d) 0.5	
	10	+5	18.	At what temperature in the	celsius scale, V (volume) of a cer	tain
	(c) $\frac{18^2}{45^2 \times 32}$	(d) $\frac{45^2}{18^2 \times 32}$		mass of gas at $27^{o}C$ we constant	vill be doubled keeping the pres	sure
10.	The ratio of rates of diffusion	1 of SO_2, O_2 and CH_4 is		() 540 5		
				(a) $54^{\circ}C$	(b) $327^{\circ}C$	
		[610 1992]		(c) $427^{o}C$	(d) $527^{o}C$	
	(a) $1:\sqrt{2}:2$	(b) $1:2:4$	19.	Pressure of a mixture of 4 g	of O_2 and 2 g of H_2 confined	in a
	(c) $2:\sqrt{2}:1$	(d) $1:2:\sqrt{2}$		bulb of 1 <i>litre</i> at $0^{o}C$ is	[AIIMS 2	000]
				(a) 25.215 <i>atm</i>	(b) 31.205 <i>atm</i>	

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	(c) 45.215 <i>atm</i>	(d) 15.210 <i>atm</i>
20.	If pressure becomes double at the	ne same absolute temperature on 2
	${\it LCO}_2$, then the volume of ${\it CO}_2$	becomes
		[A11MS 1992]
	(a) 2 <i>L</i>	(b) 4 <i>L</i>
	(c) 25 <i>L</i>	(d) 1 <i>L</i>
21.	Volume of the air that will be e	xpelled from a vessel of 300 cm^3
	when it is heated from $27^{o}C$ t	o $37^{o}C$ at the same pressure will
	be	
	(a) 310 cm^3	(b) 290 <i>cm</i> ³
	(c) 10 cm^3	(d) 37 cm ³

22. 300 *ml* of a gas at $27^{\circ}C$ is cooled to $-3^{\circ}C$ at constant pressure, the final volume is

[NCERT 1981, MP PMT 1992]

- (a) 540 *ml* (b) 135 *ml* (c) 270 *ml* (d) 350 *ml*
- Assertion & Reason

For AIIMS Aspirants

Read the assertion and reason carefully to mark the correct option out of the options given below :

- (a) If both assertion and reason are true and the reason is the correct explanation of the assertion.
- (b) If both assertion and reason are true but reason is not the correct explanation of the assertion.
- (c) If assertion is true but reason is false.
- (d) If the assertion and reason both are false.
- (e) If assertion is false but reason is true.

1.	Assertion :	Plot of P Vs. $1/V$ (volume) is a straight line.
	Reason :	Pressure is directly proportional to volume.
2.	Assertion :	Jet aeroplane flying at high altitude need pressurization of the cabin.
	Reason :	Oxygen is not present at higher altitude.
3.	Assertion :	1 mol of \boldsymbol{H}_2 and \boldsymbol{O}_2 each occupy 22.4 L of
		volume at $0^{o}C$ and 1 bar pressure.
	Reason :	Molar volume for all gases at the same temperautre and pressure has the same volume.
4.	Assertion :	Pressure exerted by a mixture of reacting gases is equal to the sum of their partial pressures.
	Reason :	Reacting gases react to form a new gas having pressure equal to the sum of both.
5.	Assertion :	Greater the value of Vander Waal's constant $'a'$ greater is the liquefaction of gas.
	Reason :	' <i>a</i> ' indirectly measures the magnitude of attractive forces between the molecules.
6.	Assertion :	Carbondioxide has greater value of root mean square velocity μ_{ms} than carbon monoxide.
	Reason :	$\mu_{\it ms}$ is directly proportional to molar mass.
7.	Assertion :	4.58 mm and $0.0098^{\circ}C$ is known to be triple point of water.
	Reason :	At this pressure and temperature all the three states i.e., water, ice and vapour exist simultaneously.

8.	Assertion	:	$1/4^{\scriptscriptstyle 6}$ of the gas is expelled if air present in an
			open vessel is heated from $27^{\circ}C$ to $127^{\circ}C$.
	Reason	:	Rate of diffusion of a gas is inversely proportional to the square root of its molecular mass.
9.	Assertion	:	Compressibility factor for hydrogen varies with pressure with positive slope at all pressures.
	Reason	:	Even at low pressures, repulsive forces dominate hydrogen gas. [AIIMS 2005]
10.	Assertion	:	vander Waal's equation is applicable only to non- ideal gases.
	Reason	:	Ideal gases obey the equation $PV = nRT$.
11.	Assertion	:	Pressure exerted by gas in a container with increasing temperature of the gas.
	Reason	:	With the rise in temperature, the average speed of gas molecules increases.
			[AIIMS 1995]
12.	Assertion	:	Gases do not settle to the bottom of container.
	Reason	:	Gases have high kinetic energy.
			[AIIMS 1997]
13.	Assertion	:	A mixture of He and O_2 is used for respiration for deep sea divers.
	Reason	:	He is soluble in blood. [AIIMS 1998]
14.	Assertion	:	Wet air is heavier than dry air.
	Reason	:	The density of dry air is more than density of water. [AIIMS 1999]
15.	Assertion	:	All molecules in a gas have some speed.
	Reason	:	Gas contains molecules of different size and shape. [A11MS 2001]
16.	Assertion	:	Effusion rate of oxygen is smaller than nitrogen.
	Reason	:	Molecular size of nitrogen is smaller than oxygen.[AIIMS 200

Answers

Characteristics and Measurable properties of gases

1	С	2	С	3	b	4	а	5	C
6	d	7	а	8	b	9	С	10	а
11	а	12	b	13	а	14	d	15	C
16	е								

Ideal gas equation and Related gas laws

1	с	2	d	3	a	4	a	5	a
6	bc	7	a	8	a	9	C	10	d
11	а	12	а	13	а	14	а	15	с
16	с	17	b	18	С	19	a	20	b
21	с	22	С	23	d	24	a	25	с
26	с	27	b	28	c	29	с	30	С

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31	a	32	а	33	b	34	b	35	b
36	C	37	d	38	c	39	c	40	a
41	b	42	с	43	c	44	d	45	a
46	a	47	b	48	d	49	b	50	c
51	d	52	а	53	а	54	c	55	a
56	а	57	b	58	а	59	b	60	a
61	b	62	d	63	c	64	b	65	d
66	b	67	d	68	а	69	а	70	b
71	с	72	а	73	d	74	c	75	a
76	с	77	d	78	b	79	b	80	b
81	d	82	d	83	С	84	C	85	a
86	d	87	а	88	а	89	C	90	c
91	a	92	C	93	а	94	b	95	a
96	а	97	d	98	b	99	е	100	а
101	с	102	с	103	с	104	с	105	a
106	а								

Kinetic molecular theory of gases and Molecular collisions

1	d	2	c	3	d	4	d	5	b
6	а	7	a	8	С	9	b	10	b
11	а	12	d	13	а	14	а	15	acd
16	а	17	b	18	d	19	b	20	d
21	a	22	d	23	с	24	b	25	b
26	a	27	d	28	а	29	d	30	c
31	C								

Molecular speeds

1	a	2	c	3	d	4	d	5	d
6	C	7	b	8	b	9	d	10	b
11	d	12	C	13	а	14	d	15	d
16	c	17	d	18	d	19	b	20	b
21	C	22	a	23	с	24	b	25	C
26	a	27	d						

Real gases and Vander waal's equation

1	b	2	b	3	c	4	c	5	d
6	C	7	d	8	b	9	d	10	b
11	C	12	b	13	b	14	d	15	c
16	b	17	а	18	b	19	C	20	a

-			_			
21	b	22	а			





Critical state and Liquefaction of gases											
1	а	2	b	3	d	4	с	5	d		
6	d	7	с	8	с	9	abc	10	c		
11	C										

Critical Thinking Questions

1	а	2	C	3	C	4	C	5	а
6	d	7	b	8	b	9	а	10	а
11	a	12	b	13	b	14	C	15	b
16	a	17	b	18	b	19	а	20	d
21	C	22	C						

Assertion & Reason

1	C	2	C	3	а	4	d	5	а
6	d	7	a	8	b	9	a	10	b
11	а	12	а	13	C	14	е	15	d
16	C								

Answers and Solutions

Characteristics and Measurable properties of gases

- 1. (c) Gases do not have definite shape and volume. Their volume is equal to the volume of the container.
- **2.** (c) All the three phases of water can coexist at $0^{o}C$ & 4.7 mm pressure.
- **3.** (b) It is characteristic of gases *i.e.* Thermal energy >> molecular attraction
- (a) In gases, molecular attraction is very less and intermolecular spaces are large hence kinetic energy of gases is highest.
 - (c) Gases and liquids, both can flow and posses viscosity.
- 7. (a) Newton is unit of force.

8. (b)
$$\frac{C^o}{5} = \frac{F^o - 32}{9}$$

5

- 9. (c) $1L = 10^{-3}m^3 = 10^3 cm^3 = 1dm^3 = 10^3 ml$.
- 10. (a) $1 atm = 10^6$ dynes cm⁴
- 12. (b) Barometer is used to measure atmospheric pressure of mixture of gases. Staglometer is used to measure surface tension. Only manometer is used to measure pressure of pure gas in a vessel.
- **13.** (a) $0^{o}C$ is equivalent to $273^{o}K$ *i.e.* conditions are same so volume will be V ml.
- 14. (d) The mass of gas can be determined by weighing the container, filled with gas and again weighing this container after removing the gas. The difference between the two weights gives the mass of the gas.
- 15. (c) Nobel gases has no intermolecular forces due to inertness.
- **16.** (e) Total volume of two flasks = 1+3=4
 - If P_1 the pressure of gas N_2 in the mixture of N_2 and O_2 then

$$P = 100 \ kPa \ , P_1 = ? \ , V = 1 \ litre \ , V_1 = 4 \ litre$$
applying Boyle's law $PV = P_1V_1$

$$100 \times 1 = P_1 \times 4 \ ; P_1 = 25$$
If P_2 is the pressure of O_2 gas in the mixture of O_2 and N_2 then, $320 \times 3 = P_2 \times 4 \ ; P_2 = 240$
Hence, Total pressure $P = P_1 + P_2 = 25 + 240$

$$= 265 \ kPa$$

Ideal gas equation and Related gas laws

(c) Boyle's law is $V \propto \frac{1}{P}$ at constant T

1.

2.

3.

4.

7.

11.

13.

(d) According to Boyle's law
$$V \propto \frac{1}{P}$$

$$V = \frac{\text{Constant}}{P}$$
; $VP = \text{Constant}$.

- (a) At sea level, because of compression by air above the proximal layer of air, pressure increases hence volume decreases *i.e.* density increases. It is Boyle's law.
- (a) At constant $T, P_1V_1 = P_2V_2$ $1 \times 20 = P_2 \times 50$; $P_2 = \frac{20}{50} \times 1$
- **5.** (a) *P.V* = constant at constant temperature. As temperature changes, the value of constant also changes.
- **6.** (b,c)According to Boyle's Law PV = constant, at constant temperature either *P* increases or *V* increases both (*b*) & (*c*) may be correct.

(a)
$$\frac{d_1}{d_2} = \frac{1}{2}$$
, $\frac{T_1}{T_2} = \frac{2}{1}$, $\frac{P_1}{P_2} = \frac{V_2}{V_1} \times \frac{T_1}{T_2} = \frac{T_1 \cdot d_1}{T_2 \cdot d_2}$
 $\frac{P_1}{P_2} = \frac{2}{1} \cdot \frac{1}{2} = \frac{1}{1}$

8. (a) Absolute temperature is temperature measured in ${}^{o}Kelvin$, expressed by T

(a)
$$T_1 = 273^\circ C = 273 + 273^\circ K = 546^\circ K$$

$$T_{2} = 0^{o} C = 273 + 0^{o} C = 273^{o} K$$

$$P_{1} = 1 ; P_{2} = ?$$
According to Gay-Lussac's law
$$\frac{P_{1}}{T_{1}} = \frac{P_{2}}{T_{2}} : P_{2} = \frac{P_{1}T_{2}}{T_{1}} = \frac{1 \times 273^{o} K}{546^{o} K} \text{ atm; } \frac{1}{2} \text{ atm.}$$

12. (a) $V_t = V_o(1 + \alpha_v t)$ $\therefore (V_2 - V_1) = \Delta V = V_o \alpha (t_2 - t_1)$ if $t_2 - t_1 = 1^o$ then $\Delta V = \alpha V_o$

For every $1^{o}C$ increase in temperature, the volume of a given mass of an ideal gas increases by a definite fraction $\frac{1}{273.15}$ of V_{o} . Here V_{o} is volume at $0^{o}C$ temperature.

(a)
$$\frac{V_1}{V_2} = \frac{T_1}{T_2}$$
 \therefore $V_2 = \frac{T_2}{T_1} V_1 = \frac{546^{\circ} K}{273^{\circ} K} \times 0.2L = 0.4L.$

4. (a)
$$V_2 = \frac{T_2}{T_1} \cdot V_1 = \frac{270^{\circ} K}{300^{\circ} K} \cdot 400 cm^3 = 360 cm^3$$

contraction
$$= V_1 - V_2 = 400 - 360 = 40 cm^3$$

15. (c) At constant volumes $P \propto T$

$$P = \text{constant } T; \quad PV = nRT \therefore P = \frac{nR}{V}T$$

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slope =
$$m = \frac{nR}{V}$$
 :: $V_2 < V_1$

 $\frac{m_1}{m_2} = \frac{V_2}{V_1} \therefore m_1 < m_2 ~~{\rm is~curve~for~V}$ has a greater slope than for V

16. (c)
$$\frac{P_1}{T_1} + \frac{P_1}{T_1} = \frac{P}{T_1} + \frac{P}{T_2}$$

 $\frac{2P_1}{T_1} = P\left(\frac{T_1 + T_2}{T_1 T_2}\right); \quad \therefore P = \frac{2P_1(T_1 T_2)}{T_1(T_1 + T_2)} = \frac{2P_1 T_2}{T_1 + T_2}$

(c) At constant *V* of a definite mass 18. $\frac{P_1}{T_1} = \frac{P_2}{T_2} \therefore \frac{P_1}{P_2} = \frac{300}{600} = \frac{1}{2}$ *i.e.* pressure increases and on increasing temperature energy of molecules increases so

the rate of collisions also increases and number of moles remains constant because there is neither addition nor removal of gas in the occurring. 224 × 10[°]

$$19. (a) Avogadro number = 6.023$$

20. (b) Compressibility =
$$\frac{1}{nRT} < 1$$
 at STP (as given)
 $nRT > PV$
 $N \times 0.0821 \times 273 > 1V_m$
 $22.41 \ litres > V_m$
 $R = 0.821$
 $T = 273 \ K$
 $P = 1$
 $n = 1$

(c) The value of universal gas constant can be expressed in 21. different units and its value would depend only on the units of the measurement. \sim 22.

(c)
$$PV = nRT$$

 $R = \frac{PV}{nT}$ = litre . *atm. K^e mole*

- 23. (d) (atm. K mol) is not a unit of R
- (a) 8.31 J.K mol 24. 1 *cal* = 4.2 *J*. $\frac{8.31}{cal.K^{-1}mol^{-1}} = 1.987$ cal K mol

$$4.2 \quad \text{and} \quad \text{an$$

31. (a)
$$P = \frac{nRT}{V} = \frac{2 \times 0.0821 \times 546}{44.8 \, l} = 2 \text{ atm.}$$

32. (a)
$$\frac{P_1V_1}{n_1T_1} = \frac{P_2V_2}{n_2T_2} \therefore n_2 = \frac{P_2V_2T_1}{P_1V_1T_2}n_1$$

at STP n_1 = one mole.
 $P_2 = 1 \text{ atm.}$
 $V_2 = 22.4 \text{ lt}$
 $T_2 = 273K$
 $n_2 = \frac{1}{1} \times \frac{22.4}{22.4} \times \frac{273}{303} \times 1 = 0.9 \text{ moles}$
 $RT_1 = 0.5 \times 0.082 \times 273^{\circ} K$

33. (b)
$$V = \frac{nRT}{P} = \frac{0.5 \times 0.082 \times 273^{\circ} K}{1} = 11.2lit$$

34. (b) $V_2 = \frac{P_1 V_1 T_2}{P_2 T_1} \Rightarrow P_1 = P$; $T_1 = 273^{\circ} K$
 $P_2 = \frac{3}{2}P$; $T_2 = T_1 + \frac{T_1}{3} = \frac{4}{3} \times 273^{\circ} K$
 $V_2 = \frac{2P}{3P} \times \frac{4}{3} \times \frac{273}{273} \times 100cc = \frac{800}{9}cc = 88.888cc$
 $= 88.9 cc$

35. (b)
$$\frac{P_{1}V_{1}}{T_{1}} = \frac{P_{2}V_{2}}{T} : \frac{P_{1}V_{1}}{P_{2}V_{2}} = \frac{T_{1}}{T_{2}}$$

36. (c) $d_{a} = 2d_{b} : 2M_{a} = M_{b}$
 $PV = nRT = \frac{m}{M}RT ; P = \frac{m}{V} \cdot \frac{RT}{M} = \frac{dRT}{M}$
 $\frac{P_{a}}{P_{b}} = \frac{d_{a}}{d_{b}} \frac{M_{b}}{M_{a}} = \frac{2d_{b}}{d_{b}} \times \frac{2M_{a}}{M_{a}} = 4$
37. (d) $n \text{ of } O_{2} = \frac{16}{32} = \frac{1}{2}$
 $n \text{ of } H_{2} = \frac{3}{2}$
Total no. of moles $= \frac{3}{2} + \frac{1}{2} = 2$
 $V = \frac{nRT}{P} = \frac{2 \times .082 \times 273}{1} = 44.8 lit = 44800 ml$
38. (c) $n = \frac{PV}{RT} = \frac{m}{M}$
 $m = \frac{MPV}{RT} = \frac{34 \times 2 \times 100}{6.002 \times 2493} = 282.4 gm$
39. (c) $V_{2} = \frac{P_{1}V_{1}}{P_{1}} \frac{T_{1}}{P_{2}} = \frac{760}{600} \times \frac{546}{273} \times 273 = 691.6 ml.$
40. (a) $\frac{P_{1}V_{1}}{n_{1}T_{1}} = \frac{P_{2}V_{2}}{n_{2}T_{2}} \therefore T_{2} = \frac{P_{2}}{P_{1}} \frac{V_{2}}{V_{1}} T_{1} \frac{m_{1}}{m_{2}}$
 $= \frac{0.75}{1} \times \frac{1}{1} \times \frac{2}{1} \times 300^{\circ} K = 450^{\circ} K$
41. (b) $V_{2} = \frac{P_{1}}{P_{2}} \frac{T_{2}}{T_{1}} \cdot V_{1} = \frac{1}{0.5} \times \frac{250}{300} \times 12000 lit = 20000 lit$
42. (c) At constant pressure
 $V \propto nT \propto \frac{m}{M}T$
 $\frac{V_{1}}{V_{2}} = \frac{m_{1}T_{1}}{m_{2}} \therefore \frac{T_{1}}{T_{2}} = \frac{V_{1}}{m_{1}} \times \frac{m_{2}}{V_{2}} = \frac{d_{2}}{d_{1}} \Rightarrow \frac{300^{\circ} K}{T_{2}} = \frac{0.75d}{d}$
 $T_{2} = \frac{300}{0.75} = 400^{\circ} K$
43. (c) $\frac{P_{1}V_{1}}{T_{1}} = \frac{P_{2}V_{2}}{T_{2}} \therefore T_{2} = \frac{P_{2}V_{2}}{P_{1}V_{1}} \times T_{1} = \frac{740}{740} \times \frac{80}{100} \times 300$
 $= 240^{\circ} K = -33^{\circ} C$
47. (b) Because H_{2} & C_{1}_{2} gases may react with each other to produce HCI gas hence Dalton's law is not applicable.
48. (d) Because $HCI \& HB_{T}$ is a reacting gas mixtures.
49. (b) NH_{3} and $HCI \& \& HB_{T}$ is a reacting gas mixtures.

HBr is a reacting gas mixture to Π_3 produce $NH_4Cl \& NH_4Br$ so Dalton's law is not applicable.

50. (c) No. of moles of lighter gas
$$=\frac{m}{4}$$

No. of moles of heavier gas
$$= \frac{m}{40}$$

Total no. of moles $= \frac{m}{4} + \frac{m}{40} = \frac{11m}{40}$
Mole fraction of lighter gas $= \frac{\frac{m}{4}}{\frac{11m}{40}} = \frac{10}{11}$

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to

Partial pressure due to lighter gas = $P_o \times \frac{10}{11}$

$$=1.1\times\frac{10}{11}=1$$
atm.

(a) *m. wt.* of $NH_3 = 17$; *m.wt.* of $N_2 = 28$ 52. *m.wt.* of $CO_2 = 44$; *m.wt.* of $O_2 = 32$

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......

56.

beacuse NH_3 is lightest gas out of these gases

$$\begin{bmatrix} r & \sqrt{MolecularWeight} \end{bmatrix}$$
55. (a) $\frac{r_g}{r_{He}} = \sqrt{\frac{M_{He}}{M_g}} \therefore M_g = M_{He} \cdot \frac{r^2_{He}}{r^2_g} = \frac{4}{(1.4)^2} = \frac{4}{1.96} = 2$

$$\begin{bmatrix} Note : 1.4 = \sqrt{2} \end{bmatrix}$$
56. (a) $r_g = \frac{1}{5} \cdot r_{H_2}$

$$\frac{M_g}{M_{H_2}} = \left[\frac{r_{H_2}}{r_g}\right]^2 = (5)^2 = 25 ; M_g = 2 \times 25 = 50$$

57. (b)
$$r_g = \frac{1}{6} r_{H_2}; \quad M_g = M_{H_2} \cdot \left[\frac{r_{H_2}}{r_g} \right]^2 = 2 \times 6^2 = 2 \times 36 = 72$$

58. (a)
$$M_1 = 64$$
; $r_2 = 2r_1$
 $M_2 = M_1 \left[\frac{r_1}{r_2} \right]^2 = 64 \times \frac{1}{4} = 16$
59. (b) $r_0 = r_H \sqrt{\frac{d_H}{d_0}} = 1 \sqrt{\frac{0.09}{1.44}} = \sqrt{\frac{1}{16}} = \frac{1}{4}$
60. (a) $r_a = 5r_b$; $\frac{d_a}{d_b} = \left[\frac{r_b}{r_a} \right]^2 = \left(\frac{1}{5} \right)^2 = \frac{1}{25}$

61. (b)
$$\frac{d_1}{d_2} = \frac{1}{16}$$
; $\frac{r_1}{r_2} = \sqrt{\frac{d_2}{d_1}} = \sqrt{16} = \frac{4}{1}$

62. (d)
$$\frac{D_A}{D_B} = \sqrt{\frac{\rho_B}{\rho_A}} = \left[\frac{\rho_B}{\rho_A}\right]^{\frac{1}{2}}; \quad \therefore D_A = D_B \left(\frac{\rho_B}{\rho_A}\right)^{\frac{1}{2}}$$

Gases may be separated by this process because of different rates of 63. (c) diffusion due to difference in their densities.

1

- (b) NH_4Cl ring will first formed near the HCl bottle because 64. rate of diffusion of $\mathit{N\!H}_3$ is more than that of HCl because $M_{NH_2}: M_{HCl} = 17:36.5$). SO NH_3 will reach first to the HCl bottle & will react there with $\,HCl$ to form $\,N\!H_4Cl$ ring
- (d) Because both $\,NO$ and $\,C_2H_6\,$ have same molecular weights 65. $\left[M_{NO} = M_{C_2H_6} = 30\right]$ and rate of diffusion \propto molecular weight. $\langle \rangle^2$

67. (d)
$$\frac{M_A}{M_B} = \left(\frac{r_B}{r_A}\right)^2$$
 :: $r_A = 2r_B$:: $\frac{r_B}{r_A} = \frac{1}{2} = \frac{1}{(2)^2} = \frac{1}{4} = .25$
68. (a) $r_H = \frac{2gm}{10}$ if $r_O = \frac{xgm}{10}$

(a)
$$r_H = \frac{r_S m}{10 \text{ min}}$$
 if $r_O = \frac{r_B m}{10 \text{ min}}$
 $r_O = r_H \sqrt{\frac{M_{H_2}}{M_{O_2}}} = \frac{2}{10} \sqrt{\frac{2}{32}}$

$$\frac{x}{10} = \frac{2}{10 \times 4} = \frac{1}{2} gm. = .5gm$$
59. (a) $r_{CH_4} = 2r_g$

$$M_g = M_{CH_4} \left(\frac{r_{CH_4}}{r_g}\right)^2 = 16 \times 2^2 = 64$$
70. (b) $r \propto \frac{1}{\sqrt{M}}$ $\therefore r = \frac{Volume \ effused}{time \ taken} = \frac{V}{t}$
 $\frac{V}{t} \propto \frac{1}{\sqrt{M}}$ \therefore for same volumes (V constant)
 $t \propto \sqrt{M}$ $\therefore \frac{t_1}{t_2} = \sqrt{\frac{M_1}{M_2}}$
 $t_{He} = t_{H_2} \sqrt{\frac{M_{He}}{M_{H_2}}} = 5\sqrt{\frac{4}{2}} = 5\sqrt{2}s.$
 $t_{O_2} = t = 5\sqrt{\frac{32}{2}} = 20s$
 $t_{CO} = 5\sqrt{\frac{28}{2}} = 5\sqrt{14}s$; $t_{CO_2} = 5\sqrt{\frac{44}{2}} = 5\sqrt{22}s$
70. (c) $\frac{r_{N_2}}{r_{SO_2}} = \frac{V_{ms}N_2}{V_{ms}SO_2} = \sqrt{\frac{T_{N_2}}{T_{SO_2}}} \cdot \frac{M_{SO_2}}{M_{N_2}} = \sqrt{\frac{T_{N_2}}{323} \times \frac{64}{28}}$
 $1.625 = \sqrt{\frac{T_{N_2}}{323} \cdot \frac{16}{7}}$
 $T_{N_2} = \frac{(1.625)^2 \times 323 \times 7}{16} = 373^o K$
72. (a) $C + H_2O \rightarrow CO_{(g)} + H_{2(g)}$
 $12gm \rightarrow 1mol + 1mol$
 $12 gm C$ produces $2mole \ or gases (1mole \ CO \ S \ 1 mole \ or \ H)$
 $\therefore 48 gm C$ may produce $\frac{48}{12} \times 2 = 4 \times 2 = 8mole$
 $= 22.4 \times 8 \ t \ gases = 179.2 \ t \ gas.$
73. (d) Molecular weight $= \frac{mRT}{PV} = \frac{4.4 \times .082 \times 273}{1 \times 2.24} = 44$

So the gas should be CO_2

74.

75.

(c)
$$PV = nRT$$

 $P = \frac{n}{V}RT$ $\therefore \frac{n}{V} = C \implies P$
 P 1

$$T = \frac{1}{CR} = \frac{1}{1 \times .821} = 12^{6} K$$
(a) 6.02×10^{22} molecules of each N_2, O_2 and H_2
 $= \frac{6.02 \times 10^{22}}{6.02 \times 10^{23}}$ moles of each
Weight of mixture = weight of 0.1 mole N_2 + weight of 0.1
mole H_2 + weight of 0.1 mole of O_2

= CRT

$$=(28 \times 0.1) + (2 \times 0.1) + (32 \times 0.1) = 6.2gm$$

76. (c) *M.wt* of
$$CO_2 = 12+16+16 = 44$$

Volume of 44 *gm* of CO_2 at NTP = 22.4 *litre*
1 *gm* of CO_2 at NTP = $\frac{22.4}{44}$



4.4 gm of
$$CO_2$$
 at N.T.P

$$\Rightarrow \frac{22.4}{44} \times 4.4 \text{ litre} = 2.24 \text{ litre}$$

78. (b) No. of moles of CO_2 present in 200 *ml* solution

= molarity × Volume (in
$$lt$$
) = $0.1 \times \frac{200}{1000} = .02$
Volume of 0.02 mole of $CO_2 = 22.4 \times .02lt$ = 0.448 lit

79. (b) Molecular weight = $V.d \times 2 = 11.2 \times 2 = 22.4$ Volume of 22.4 gm Substance of NTP = 22.4 *litre* 1 gm substance at NTP = $\frac{22.4}{22.4}$ *litre* 11.2 gm substance of NTP = 11.2 *litre*

80. (b)
$$\frac{M.wt.of O_2}{M.wt.of SO_2} \Rightarrow \frac{M_1}{M_2} \Rightarrow \frac{32}{64} = \frac{1}{2}$$

The weight of oxygen will be $\frac{1}{2}$ that of SO_2

- **81.** (b) For *HI* has the least volume because of greater molecular weight $V \propto \frac{1}{M}$
- 83. (c) Since no. of molecules is halved so pressure should also be halved.
- 84. (c) H_2 will be filled first because of lower molecular weight
- 85. (a) Mixture of SO₂ and Cl₂ are reacted chemically and forms SO₂Cl₂. That is why mixture of these gases is not applicable for Dalton's law.
 86. (d) According to Boyle's law

$$P_{1}V_{1} = P_{2}V_{2} \implies P_{1} \times 60 = 720 \times 100$$
$$P_{1} = \frac{720 \times 100}{60} = 1200mm$$

87. (a) Rate of diffusion
$$\propto \frac{1}{\sqrt{Molecular Mass}}$$

that is why H_2 gas diffuse first

88. (a) Solution level will rise, due to absorption of CO_2 by sodium hydroxide.

$$2NaOH + CO_2 \rightarrow Na_2CO_3 + H_2O$$
89. (c)
$$CaCo_3 \rightarrow CaO + CO_2 \uparrow$$

$$\therefore \text{ At S.T.P. 100g } CaCO_3 \text{ produce} = 22.4 litre \text{ of } CO_2$$

$$\therefore \text{ At S.T.P. 1g } CaCO_3 \text{ produce} = \frac{22.4}{100} = .224 litre \text{ of } CO_2$$

90. (c) The density of gas
$$= \frac{Molecularwt. Of Metal}{Volume} = \frac{45}{22.4}$$

$$= 2gmlitr\bar{e}^{-1}$$

91. (a)
$$M_1 = 32g$$
 for O_2 , $M_2 = 2g$ for H_2
 $\frac{r_1}{r_2} = \sqrt{\frac{M_2}{M_1}}$; $\frac{r_1}{r_2} = \sqrt{\frac{2}{32}} = \sqrt{\frac{1}{16}} = \frac{1}{4}$

92. (c) In 22.4*l* of
$$H_2$$
 maximum number of molecules
= 6.023×10^{23}

In 1*l* of *H*₂ maximum number of molecules = $\frac{6.023 \times 10^{23}}{22.4}$

In 15*l* of
$$H_2$$
 maximum number of molecules
= $\frac{6.023 \times 10^{23}}{22.4} \times 15 = 4.03 \times 10^{23}$ molecules.

93. (a) 22.4*l*
$$O_2$$
 at S.T.P. = 32*gm* of O_2

11
$$O_2$$
 at S.T.P. = $\frac{32}{22.4} = 1.43 gm$ of O_2

95 (a) We know that molecular mass of hydrogen $M_1 = 2$ and that of helium $M_2 = 4$, we also know that Graham's law of diffusion

$$\frac{r_1}{r_2} = \sqrt{\frac{M_2}{M_1}} = \sqrt{\frac{4}{2}} = \sqrt{2} = 1.4; r_1 = 1.4m$$
(a) $\frac{r_A}{r_H} = \sqrt{\frac{M_H}{M_A}} = \frac{r}{6r} = \sqrt{\frac{2}{M_A}}$
 $M_A = 6 \times 6 \times 2 = 72g$

97. (d) Given that:

$$V_1 = 100ml, P_1 = 720mm, V_2 = 84ml, P_2 = ?$$

By using $P_1V_1 = P_2V_2$ [According to the Boyle's law]

$$P_2 = \frac{P_1 V_1}{V_2} = \frac{720 \times 100}{84} = 857.142$$

Hence, $P_2 = 857.14mm$

96.

99.

100.

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$$PV = nRT, \ n = \frac{1}{RT}$$

$$\frac{n_A}{n_B} = \frac{\frac{P_1V_1}{RT_1}}{\frac{P_2V_2}{RT_2}}; \ \frac{n_A}{n_B} = \frac{P_1V_1}{T_1} \times \frac{T_2}{P_2V_2}$$

$$\frac{n_A}{n_B} = \frac{2P \times 2V}{2T} \times \frac{T}{PV}; \ \frac{n_A}{n_B} = \frac{2}{1}$$
(e) No. of molecules $= 2 \times V.d$
 $2 \times 38.3 = 76.3$
wt. of $NO_2 = x$
So that wt. of $N_2O_4 = 100 - x$
Hence, $\frac{x}{46} + \frac{100 - x}{92} = \frac{100}{76.6} = \frac{2x + 100 - x}{92} = \frac{100}{76.6}$
 $x = 20.10$, no. of mole. of $NO_2 = \frac{20.10}{46} = 0.437$
(a) Given that
 $P_1 = 76cm$ of Hg (Initial pressure at N.T.P.)

$$P_2 = ?, V_1 = 5litre, V_2 = 30 + 5 = 35litres$$

According to Boyle's law
$$P_1V_1 = P_2V_2; 76 \times 5 = P_2 \times 35$$

$$P_2 = \frac{76 \times 5}{35} \implies P_2 = 10.8cm \text{ of } Hg$$

101. (c) Given initial volume $(V_1) = 300cc$, initial temperature $(T_1) = 27^{\circ}C = 300K$, initial pressure $(P_1) = 620mm$, final

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temperature
$$T_2 = 47^{\circ}C = 320K$$
 and final pressure
 $(P_2) = 640mm$. We know from the general gas equation
 $\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2} = \frac{620 \times 300}{300} = \frac{640 \times V_2}{320} \Rightarrow V_2 = 310cc$
102. (c) $NH_3 + HCl \rightarrow NH_4Cl$
 $4litre 1.5litre$
 HCl is a limiting compound. That's why 1.5litre of HCl
reacts with 1.5litre of NH_3 and forms NH_4Cl . Thus (4 -
1.5) 2.5litre NH_3 remains after the reaction.
103. (c) $\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$; $\frac{P_1 \times 4}{T_1} = \frac{2P_1 \times V_2}{2T_1}$
 $8 = 2 \times V_2$ so $V_2 = 4dm^3$
104. (c) $P_1 = P, V_1 = V, T_1 = T$
 $P_2 \frac{P}{2}, V_2 = ?, T_2 = T$
According to gas equation
 $\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$ or $\frac{PV}{T} = \frac{PV_2}{2T}$
 $\therefore V_2 = 4V$
105. (a) $P_1 = P, V_1 = V, T_1 = 273 + 75 = 348K$
 $P_2 = 2P, V_2 = \frac{85}{100}, T_2 = ?$
 $\frac{P \times V}{398} = \frac{2P \times 85V}{T_2 \times 100} \Rightarrow T_2 = \frac{348 \times 2 \times 85}{100}$
 $\therefore T_2 = 591.6K = 318.6^{\circ}C$
106. (a) Boyle's law $-PV =$ constant
On differentiating the equation,
 $d(PV) = d(C) \Rightarrow PdV + VdP = 0$
 $\Rightarrow VdP = -PdV \Rightarrow \frac{dP}{P} = -\frac{dV}{V}$.

Kinetic molecular theory of gases and Molecular collisions

4. (d) Kinetic energy
$$=\frac{3}{2}RT$$

- 5. (b) $K.E = \frac{3}{2}RT$ it means that *K.E* depends upon *T* (absolute temperature) only.
- 7. (a) $KE = \frac{3}{2}RT = \frac{3}{2}PV$ $\therefore P = \frac{2}{3}\frac{E}{V}$ for unit volume $(V = 1) \implies P = \frac{2}{3}E$
- 8. (c) Tr. $K.E. = \frac{3RT}{2}$ it means that the Translational Kinetic energy of Ideal gas depends upon temperature only.
- 9. (b) $\frac{E_{He}}{E_{H_2}} = \sqrt{\frac{T_{He}}{T_{H_2}}}$ so energies will be same for *He* & *H*₂ at same temperature.

n. (a)
$$K.E. = \frac{3}{2}.RT = \frac{3}{2}.2.T$$
 $\therefore R \approx 2calK^{-1}mol^{-1}$
 $K.E. = 3T$

- 12. (d) All molecules of an ideal gas show random motion. They collide with each other and walls of container during which they lose or gain energy so they may not have same kinetic energy always.
- 13. (a) For same temperature kinetic energies of H_2 & He molecules will be same because kinetic energy depends only on temperature.
- 14. (a) For same temp. kinetic energies would be equal for all molecules, what ever their molecular weights will be, it doesn't matter.
- (a,c,d)Kinetic energies per molecule will be same because it is proportional to absolute temperature only.

$$\frac{d_{N_2}}{d_{CO_2}} = \frac{M_{N_2}}{M_{CO_2}} = \frac{28}{44} \text{ i.e. } dN_2 < dCO_2$$

Total translational kinetic energy will also be same because at same temperature & pressure number of molecules present in same volume would be same (according to Avogadro's Law)

- (a) On increasing pressure, the volume decreases and density increases. So molecules get closer to each other hence mean free path also decreases.
- 17. (b) Most probable velocity increase and fraction of molecule possessing most probable velocity decrease.

18. (d)
$$PV = nRT = \frac{m}{M}RT$$

 $\therefore \frac{m}{V} = \frac{PM}{RT} = \text{density}$

16.

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9. (b)
$$d \propto \frac{P}{T}$$
 the value of $\frac{P}{T}$ is maximum for (b)

20. (d) If inversion temperature is $80^{\circ}C = 193^{\circ}K$ then the temperature, at which it will produce cooling under Joule Thomson's effect, would be below inversion temperature except $173^{\circ}K$ all other values given as

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21. (a) Since
$$\frac{C_P}{C_V} = 1.4$$
, the gas should be diatomic.

If volume is 11.2 *lt* then, no. of moles =
$$\frac{1}{2}$$

 \therefore no. of molecules = $\frac{1}{2} \times$ Avagadro's No.

2

no. of atoms =
$$2 \times no.$$
 of molecule

$$2 \times \frac{1}{2} \times \text{Avagadro's No.}$$

$$= 6.0223 \times 10^{23}$$

1

22. (d) Density = $\frac{M}{V}$ $d = \frac{v.d \times 2}{V}$

$$V = \frac{V}{2}$$

$$V.d = \frac{d \times V}{2}$$

$$V.d = \frac{0.00130 \times 22400}{2} = 14.56 gm^{-1}$$

 $(M = V.d \times 2)$

23. (c) Volume of steam =
$$1t = 10^3 cm^3$$

 $\therefore m = d.V$
 $\therefore mass of 10^3 cm^3 steam = density \times Volume$
 $= \frac{0.0006gm}{cm^3} \times 10^3 cm^3 = 0.6gm$

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Actual volume occupied by $H_2 {\cal O}\,$ molecules is equal to volume of water of same mass

- \therefore Actual volume of $\,H_2O\,$ molecules in $\,6\,gm\,$ steam $\,$
- = mass of steam/density of water

$$= 0.6 gm / 1 gm/cm \Rightarrow 0.6 cm^3$$

24. (b)
$$r = \frac{C_P}{C_V} = \frac{5}{3} = 1.66$$
 (For Monoatomic as He, Ne, Ar)

25. (b) The density of neon will be highest at $0^{\circ}C$ 2 *atm* according to $d \propto \frac{P}{T}$

29. (d) *K.E.* per mole
$$=\frac{3}{2}RT$$

so all will have same K.E. at same temperature.

30. (c)
$$\because W = P.dV = E$$

 \therefore Energy per unit volume = P
31. (c) $d \propto M \implies \frac{d_1}{d_2} = \frac{M_1}{M_2}; \frac{3d}{d} = \frac{M}{M_2}; M_2 = \frac{M}{3}.$

Molecular speeds

1. (a)
$$V_{ms} = \sqrt{\frac{3RT}{M}}, V_{av} = \sqrt{\frac{8RT}{\pi M}}; \frac{V_{ms}}{V_{av}} = \sqrt{\frac{3\pi}{8}}$$
$$= \sqrt{\frac{66}{56}} \Rightarrow \frac{1.086}{1}$$

 $\textbf{3.} \qquad (d) \quad most \ probable \ velocity: mean \ velocity: V_{_}$

$$= \sqrt{\frac{2RT}{M}} : \sqrt{\frac{8RT}{\pi M}} : \sqrt{\frac{3RT}{M}} = \sqrt{2} : \sqrt{\frac{8}{\pi}} : \sqrt{3}$$
4. (d) $V_{ms} = \sqrt{\frac{3RT}{M}} \therefore V_{ms} \propto \frac{1}{\sqrt{M}}$ at same T

because H_2 has least molecular weight so its r.m.s. velocity should be maximum.

5. (d)
$$\frac{U_{SO_2}}{U_{He}} = \frac{1}{2} = \sqrt{\frac{M_{He}}{M_{SO_2}}} \frac{T_{SO_2}}{T_{He}} = \sqrt{\frac{4}{64}} \cdot \frac{T_{SO_2}}{300}$$

 $= \frac{4}{64} \cdot \frac{T_{SO_2}}{300} = \frac{1}{4}; \quad T_{SO_2} = 1200^o K$
6. (c) $\frac{U_{O_3}}{U_{O_2}} = \sqrt{\frac{M_{O_2}}{M_{O_3}}} = \sqrt{\frac{32}{48}} = \sqrt{\frac{2}{3}}$

7. (b) Average kinetic energy per molecule

$$=\frac{3}{2}KT = \frac{3}{2} \times 1.38 \times 10^{-23} \times 300J = 6.17 \times 10^{-21}J$$

8. (b)
$$\frac{U_{SO_2}}{U_{O_2}} = \sqrt{\frac{M_{O_2}T_{SO_2}}{M_{SO_2}T_{O_2}}} = \sqrt{\frac{32 \times T_{SO_2}}{64 \times 303}} = 1$$

 $1 = \frac{32 \times T_{SO_2}}{64 \times 303} \Rightarrow T_{SO_2} = 606$

9. (d) Among these Cl_2 has the highest molecular weight so it will posses lowest root mean square velocity.

10. (b)
$$\frac{U_1}{U_2} = \sqrt{\frac{T_1}{T_2}}$$
 $\therefore \frac{T_1}{T_2} = \left(\frac{5 \times 10^4}{10 \times 10^4}\right)^2 = \frac{1}{4}$

12. (c)
$$V_{ms} = \sqrt{\frac{3KT}{Molecularweight}}$$
 i.e. $V_{ms} \propto \frac{1}{\sqrt{m}} \propto (m)^{-\frac{1}{2}}$

 (a) When average speed of molecule is increased due to increase in temperature then the change in momentum during collision between wall of container and molecules of gas also increases.

14. (d)
$$\frac{U_1}{U_2} = \sqrt{\frac{m_2}{m_1}} \cdot \frac{T_1}{T_2} \quad \because T_1 = T_2$$

 $\frac{U_1^2}{U_2^2} = \frac{m_2}{m_1} \qquad \because m_1 U_1^2 = m_2 U_2^2$
15. (d) $U_2 = U_1 \sqrt{\frac{T_2}{T_1}} = U_1 \sqrt{\frac{1200}{300}} = U_1 \times 2$

r.m.s. velocity will be doubled.

16. (c)
$$\frac{U_{H_2}}{U_{O_2}} = \sqrt{\frac{T_{H_2}}{M_{H_2}}} \cdot \frac{M_{O_2}}{T_{H_2}} = \sqrt{\frac{50}{2}} \cdot \frac{32}{800} = 1$$

17. (d)
$$U = \sqrt{\frac{3RT}{M}} = \sqrt{\frac{3PV}{M}} = \sqrt{\frac{3P}{d}} \therefore U \propto \frac{1}{\sqrt{d}}$$

19. (b)
$$\frac{U_1}{U_2} = \sqrt{\frac{n_1 T_1}{n_2 T_2}} = \sqrt{\frac{n \times T}{2n \times 2T}} = \sqrt{\frac{1}{4}} = \frac{1}{2}$$

 $U_2 = 2U_1 = 2U$

21. (c)
$$\frac{V_{mp}}{V_{av}} = \frac{\sqrt{\frac{2RT}{M}}}{\sqrt{\frac{8RT}{\pi M}}} = \frac{\sqrt{\pi}}{2}$$
22. (a)
$$V_{ms} = \sqrt{\frac{3RT}{M}} \Rightarrow V_{ms} = \sqrt{T}$$
Given $V_1 = V$, $T_1 = 300K$, $V_2 = 2V$, $T_2 = ?$

26.

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$$\frac{V_1}{V_2} = \sqrt{\frac{T_1}{T_2}} : \left(\frac{V}{2V}\right)^2 = \frac{300}{T_2} \implies T_2 = 300 \times 4 = 1200K$$
24. (b) $\left(P - \frac{n^2 a}{V^2}\right) (V - nb) = nRT$
 $\left(P - \frac{(2)^2 \times 4.17}{(5)^2}\right) (5 - 2 \times .03711) = 2 \times .0821 \times 300$

$$P = \frac{2 \times .0821 \times 300}{5 - 2 \times .03711} - \frac{4.7 \times 2^2}{5^2} \Longrightarrow 10 - 0.66 = 9.33 atm}$$
(a) Average speed : most probable speed

$$\sqrt{\frac{8RT}{\pi M}} : \sqrt{\frac{2RT}{M}} \Rightarrow \sqrt{\frac{8}{\pi}} : \sqrt{2} \Rightarrow 1.128 : 1.$$

27. (d)
$$v_{mms} = \sqrt{\frac{3RT}{M}}$$

 $\sqrt{30^2 R} = \sqrt{\frac{3RT}{M}} \Rightarrow 30 \times 30R = \frac{3R \times 300}{M}$
 $\Rightarrow M = \frac{3 \times 300}{30 \times 30} 1 gm = 0.001 kg$.

Real gases and Vander waal's equation

2. (b) Because molecules of real gases have intermolecular forces of attraction so the effective impact on the wall of container is

diminished. Pressure of real gas is reduced by $\frac{a}{v^2}$ factor hence

behaviour of real gas deviate from ideal behaviour. עת

3. (c)
$$Z = \frac{PV}{RT}$$
 : for ideal gas $PV = RT$ so $Z = I$
12. (b) $Z = \frac{PV}{RT}$; for ideal gas $PV = RT$; so $Z = I$

- 13. (b) Ideal gas has no attractive force between the particles
- PV = nRT is a ideal gas equation it is allowed when the 14. (d) temperature is high and pressure is low.
- 16. (b) At Boyle temperature real gas is changed into ideal gas When pressure is low (a) 17.

$$\begin{bmatrix} P + \frac{a}{V^2} \end{bmatrix} (V - b) = RT$$

or $PV = RT + Pb - \frac{a}{V} + \frac{ab}{V^2}$ or $\frac{PV}{RT} = 1 - \frac{a}{VRT}$
 $Z = -\frac{a}{VRT} \left(\because \frac{PV}{RT} = Z \right)$

- 18. (b) At high temperature and low pressure, Vander Waal's equation is reduced to ideal gas equation. PV = nRT
 - PV = RT (For 1 mole of gas)
- Vander waal's constant for volume correction b is the measure 20. (a) of the effective volume occupied by the gas molecule.

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

 $\left(P + \frac{2.253}{0.25 \times 0.25}\right)(0.25 - 0.0428) = 0.0821 \times 300$
or $(P + 36.048)(0.2072) = 24.63$

 \Rightarrow P+36.048 = 118.87 \Rightarrow P = 82.82 atm.

Critical state and Liquefaction of gases

- A diabatic demagnetisation is a technique of liquefaction of 2. (b) gases in which temperature is reduced.
- (d) An ideal gas can't be liquefied because molecules of ideal gas 3. have not force of attraction between them.
- (c) At above critical temperature, substances are existing in 4 gaseous state, since gas cannot be liquefied above it.
- 5 (d) Absence of inter molecular attraction ideal gas cannot be liquefied at any volume of *P* and *T*. 6.
 - (d) For Z gas of given gases, critical temperature is highest

$$T_c = \frac{8a}{27Rb} \Rightarrow T_c = \frac{8 \times 12}{27 \times .0821 \times .027} = 1603.98K$$

- (c) Value of constant a is greater than other for NH_3 that's why 7. NH_3 can be most easily liquefied.
- 8. The temperature below which the gas can be liquefied by the (c) application of pressure alone is called critical temperature.

10. (c)
$$d = \frac{M}{V} \implies M = d \times V$$

 $M = 2.68 \times 22.4$ at N.T.P. (:: $V = 22.41$)
 $M = 60.03 gm$
m. wt of $COS = 12 + 16 + 32 = 60$

11. (c) 22400 *ml* is the volume of
$$O_2$$
 at N.T.P =32*gm* of O_2

1m is the volume of O_2 at NTP = $\frac{32}{22400}$ 112 *ml* is the volume of O_2 at NTP = $\frac{32}{22400} \times 112$ = 0.16 gm of O_2

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Critical Thinking Questions
1. (a) Average kinetic energy
$$\propto (T \text{ Kelvin})$$

(Factor) $\frac{K.E_2}{K.E_1} = \frac{T_2}{T_1} = \frac{40 + 273}{20 + 273} = \frac{313}{293}$
2. (c) *M. wt.* = *V.d.* × 2
 $= 70 \times 2 = 140 \implies x = \frac{m.wt.}{wt.of[CO]} = \frac{140}{[12+16]} = 5$
3. (c) Gas deviate from ideal gas behaviour to real gas (according to Vander Waal's at low temperature and high pressure)

(c) At same temperature and pressure, equal volumes have equal number of molecules. If 1/it. of oxygen consists N molecules then at same temperature and pressure 1 lit of SO_2 will consists N molecules. So 2 lit. of SO_2 will contain 2Nmolecules.

(a)
$$V_{av}: V_{ms}: V_{ms} = V: U: \alpha$$

 $\sqrt{\frac{8RT}{\pi M}}: \sqrt{\frac{3RT}{M}}: \sqrt{\frac{2RT}{M}}$
 $\alpha: V: U = \sqrt{2}: \sqrt{\frac{8}{\pi}}: \sqrt{3} = 1:1.128:1.224$

6. (d)
$$\frac{C_p}{C_v}$$
 ratio for diatomic gases is 1.40 at lower temperature &

(b)
$$PV = nRT$$
 (For ideal gas)

$$Z = \frac{PV}{nRT} = 1$$
 (For ideal gas)

(b) If
$$Z < 1$$
 then molar volume is less than 22.4 L

9. (a)
$$r_{O_2} = \frac{x}{18} mole / \sec \Rightarrow r_g = \frac{x}{45} mol / \sec$$

$$M_g = M_{O2} \left(\frac{r_{O_2}}{r_g}\right)^2 = 32 \left(\frac{x}{18} \times \frac{45}{x}\right)^2 = 32 \times \frac{45^2}{18^2}$$

$$\begin{array}{ll} \textbf{10.} & \textbf{(a)} & r_{SO_2} : r_{CH_4} = \frac{1}{\sqrt{M_{SO_2}}} : \frac{1}{\sqrt{O_2}} : \frac{1}{\sqrt{CH_4}} \\ & = \frac{1}{\sqrt{64:32:16}} = \frac{1}{\sqrt{4:2:1}} \\ & \frac{1}{2} : \frac{1}{\sqrt{2}} : \frac{1}{1}; \quad \frac{2}{2} : \frac{2}{\sqrt{2}} : \frac{2}{1}; \quad 1 : \sqrt{2} : 2 \end{array}$$

I. (a) Root mean square speed =
$$\begin{bmatrix} n_1c_1^2 + n_2c_2^2 + n_3c_3^2 + \dots \\ n_1 + n_2 + n_3 + \dots \\ n_1 + n_2 + n_3 + \dots \end{bmatrix}$$

12. (b) If 40 m/
$$O_2$$
 will diffuse in t min. then. $r_{O_2} = \frac{40}{t}$

$$r_{H} = \frac{50}{20} \implies r_{O} = r_{H} 2\sqrt{\frac{M_{H_{2}}}{M_{O_{2}}}} = \frac{50}{20}\sqrt{\frac{2}{32}} = \frac{50}{20} \cdot \frac{1}{4}$$
$$\frac{40}{t} = \frac{50}{80} \therefore t = \frac{40 \times 80}{50} = 60 \text{ min.}$$

13. (b)
$$\frac{V_{av}CH_4}{V_{ab}O_2} = \sqrt{\frac{T_{CH_4}}{T_{O_2}} \cdot \frac{M_{O_2}}{M_{CH_4}}} = 1$$

 $\frac{T_{CH_4}}{300} \cdot \frac{32}{16} = 1$; $T_{CH_4} = 150^{\circ} K$

(c) Pressure of O_2 (dry) = 751-21 = 730 mm Hg 14.

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$$= \frac{730}{760} = 0.96atm$$
15. (b) $PV = nRT$, $n = \frac{PV}{RT} = \frac{1 \times 145}{0.082 \times 3} = 5.8 \approx 6 \text{ mole}$.
16. (a) $V = \frac{nRT}{P} = \frac{m}{M} \cdot \frac{RT}{P} = \frac{1}{58} \times \frac{0.82 \times 350}{1} = 0.495lit$
 $= 495 \text{ cm}$
17. (b) $\frac{r_{He}}{r_{CH_4}} = \sqrt{\frac{M_{CH_4}}{M_{He}}} = \sqrt{\frac{16}{4}} = 2$
18. (b) $\frac{V_1}{V_2} = \frac{T_1}{T_2} \because T_2 = \frac{T_1V_2}{V_1} = 300^{\circ} K, \frac{2V}{V} = 600^{\circ} K$
 $T_2 = 600^{\circ} K = (600 - 273)^{\circ} C = 327^{\circ} C$
19. (a) no. of moles of $O_2 = \frac{4}{32} = 0.125$
no. of moles of $H_2 = \frac{2}{2} = 1$
total no. of moles $1 + 0.125 = 1.125$
 $P = \frac{nRT}{V} = \frac{1.125 \times 0.082 \times 273}{1} = 25.184atm$.
20. (d) $\frac{P_1}{P_2} = \frac{1}{2}, \because \frac{V_1}{V_2} = \frac{P_2}{P_1} = \frac{2}{1}$
 $\frac{2L}{V_2} = \frac{2}{1}; V_2 = 1L$
21. (c) $\frac{V_2}{V_1} = \frac{T_2}{T_1}$
 $\therefore V_2 = \frac{T_2}{T_1}V_1 = \frac{310^{\circ} K}{300^{\circ} K} \times 300cm^3 = 310cm^3$
22. (c) $V_2 = \frac{T_2}{T_1} \cdot V_1 = \frac{270^{\circ} K}{300^{\circ} K} \times 300ml = 270ml$

Assertion & Reason

- 1. (c) Pressure is inversly proportional to volume (Boyle's law). $p \alpha \frac{1}{V}$ (*n*, *T* constant).
- (c) The air pressure decreases with increase in altitude. So the partial pressure of Oxygen is not sufficient for breathing at higher altitude and thus pressurization is needed.
- (a) At a given temperature and pressure the volume of a gas is directly proportional to the amount of gas Van (P and T constant).
- (d) According to Dalton's law of partial pressure, the pressure exerted by a mixture of non interacting gases is equal to the sum of their partial pressures (pressure exerted by individual gases in mixture) P_{Total} = P₁ + P₂ + P₃ ... (*T* and *V* constant). Both the gases if non-interacting would spread uniformly to occupy the whole volume of the vessel.
- **5.** (a) Considering the attractive force pressure in ideal gas equation (PV = nRT) is correct by introducing a factor of $\frac{an^2}{V^2}$ where *a* is a vander waal's constant.

6. (d)
$$\mu_{ms} = \sqrt{\frac{3RT}{M}}$$
 is inversiy related to molecular mass.
Therefore, $\mu_{ms}(CO) > \mu_{ms}(CO_2)$.

8. (b) $\frac{V_1}{T_1} = \frac{V_2}{T_2}$ (Initial fraction $\frac{V_1}{V_2} = 1$ when temperature is $27^{\circ}C$. At $127^{\circ}C$ the new fraction is $\frac{V_1}{V_2} = \frac{300}{400} = \frac{3}{4}$ \therefore

r expelled
$$\Rightarrow 1 - \frac{3}{4} = \frac{1}{4}$$

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9.

- (a) In case of H, compressibility factor increases with the pressure. At 273 K, Z > 1 which shows that it is difficult to compress the gas as compared to ideal gas. In this case repulsive forces dominate.
- 10. (b) In real gases, the intermolecular forces of attraction and the volume occupied by the gas molecules cannot be neglected.
 11. (a) When the temperature increase, the average speed of gas
 - (a) When the temperature increase, the average speed of gas molecules increases and by this increase the pressure of gas is also increases.
- (a) It is correct that gases do not settle to the bottom of container and the reason for this is that due to higher kinetic energy of gaseous molecules they diffuse.
- 13. (c) The assertion, that a mixture of helium and oxygen is used for deep sea divers, is correct. The *He* is not soluble in blood. Therefore, this mixture is used.
- (e) Dry air is heavier than wet air because the density of dry air is more than water.
- 15. (d) All molecule of a gas have different speed. Therefore, they move by its own speed.
- 16. (c) Assertion is true but reason is false because of effusion rate
 - $\propto \frac{1}{\sqrt{M}}$ (Molecular weight) but it does not depend on molecular size.

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Gaseous State

ET Self Evaluation Test -6

Which of the following volume (V) – temperature (T) plots 7. Same mass of CH_4 and H_2 is taken in container. The partial 1. represents the behaviour of one mole of an ideal gas at one [IIT Screening 2002] pressure caused by H_2 is [IIT 1989; CPMT 1996] atmospheric pressure V(L)V(L)(a) 8/9 (b) 1/9 (36.8 L (28.6 L (c) 1/2 (d) 1 (22.4 L (22.4 L 373 K) 373 K 273 K) 273 K) 2. The following graph illustrates [JIPMER 2000] (a) (b) $\tau(\vec{k})$ T(K)V(L)V(L)(30.6 L (22.4 L (22.4 L 373 K) 273 K) 273 K) (14.2 L Temp. (°*C*) (d) (c) 373 K) (a) Dalton's law (b) Charle's law If the average velocity of $N_2^{\mathcal{T}(K)}$ molecule is $0.3 \, m/s$ at $2\mathcal{T}^{\mathcal{T}(K)}_{C}$ 8. (c) Boyle's law (d) Gay-Lussac's law then the velocity will be 0.6 m/s at [Pb. CET 2001] If the pressure and absolute temperature of 2 $\it litres$ of $\it CO_2$ are з. 600K(a) 1200K (b) doubled, the volume of CO_2 would become (c) 400*K* (d) 1800K [CBSE PMT 1991] Equal volumes of two gases which do not react together are 9. (a) 2 litres (b) 4 litres enclosed in separate vessels. Their pressures at 100 mm and 400 (c) 5 *litres* mm respectively. If the two vessels are joined together, then what (d) 7 litres will be the pressure of the resulting mixture (temperature remaining What is kinetic energy of 1g of O_2 at $47^{\circ}C$ 4 constant) [CBSE PMT 1981] (a) 125 mm (b) 500 mm [Orissa JEE 2004] (c) 1000 mm (d) 250 mm (a) $1.24 \times 10^2 J$ (b) $2.24 \times 10^2 J$ A gas of volume 100 cc is kept in a vessel at pressure 10^4 Pa 10. $1.24 \times 10^{3} J$ (d) $3.24 \times 10^2 J$ (c) maintained at temperature $24^{o}C$. If now the pressure is increased The root mean square speeds at STP for the gases H_2, N_2, O_2 5 to $10^5 Pa$, keeping the temperature constant, then the volume of and HBr are in the order the gas becomes [AFMC 1992] [Pb. CET 1994; CBSE PMT 1991] (a) 10 cc (b) 100 cc (c) 1 *cc* (d) 1000 cc (a) $H_2 < N_2 < O_2 < HBr$ 11. If a gas is expanded at constant temperature [IIT 1986] (b) $HBr < O_2 < N_2 < H_2$ (a) The pressure increases (b) The kinetic energy of the molecules remains the same (c) $H_2 < N_2 = O_2 < HBr$ (c) The kinetic energy of the molecules decreases (d) $HBr < O_2 < H_2 < N_2$ The number of molecules of the gas increases (d) The rate of diffusion of SO_2 and O_2 are in the ratio 12. 6. By what ratio the average velocity of the molecule in gas change when the temperature is raised from 50 to $200^{\circ} C$ [Assam JET 1991; EAMCET 1980] [DCE 2003] (a) $1:\sqrt{2}$ (b) 1:32 (a) 1.21 / 1 (b) 1.46 / 1 (c) 1:2 (d) 1:4 (c) 1.14 / 1 (d) 4 / 1







(SET -6)

1. (a)
$$N_{CH4} =$$
 number of moles of $CH_4 = \frac{m}{16}$
 $N_{H2} =$ number of moles of $H_2 = \frac{m}{2}$
fraction partial pressure of H_2 is
 $H_2 = \frac{n_{H_2}}{n_{H_2} + n_{CH_4}} = \frac{\frac{m}{2}}{\frac{m}{2} + \frac{m}{16}} = \frac{\frac{m}{2}}{\frac{9m}{16}} = \frac{8}{9}$
2. (b) According to Charle's Law $V \propto T$
 $V_t = V_o + V_o \alpha t$
compare it with $Y = C + mx$
3. (a) $V2 = \frac{P_1V_1}{T_1} \cdot \frac{T_2}{P_2} = \frac{P}{2P} \times 2lt \times \frac{2T}{T} = 2lt$
4. (a) K.E. $= \frac{3}{2}nRT = \frac{3}{2} \times \frac{1}{32} \times 8.314 \times 320J$.
 $= 1.24 \times 10^2 J$
5. (b) $V_{ms} \propto \frac{1}{\sqrt{m}}$
 $U_{H_2} : U_{N_2} : U_{O_2} : U_{HBr} = \frac{1}{\sqrt{2}} : \frac{1}{\sqrt{20}} : \frac{1}{\sqrt{32}} : \frac{1}{\sqrt{81}}$ is
 $U_{HBr} < U_{O_2} < U_{N_2} < U_{H_2}$
6. (c) $T_1 = 150 + 273 = 423K$; $T_2 = 50 + 273 = 323K$

Hence,
$$\frac{(V_{av})_1}{(V_{av})_2} = \sqrt{\frac{T_1}{T_2}} = \sqrt{\frac{423}{323}} = \frac{1.14}{1}$$

7. (c)
$$\frac{V_1}{V_2} = \frac{T_1}{T_2} \therefore V_2 = V_1 \cdot \frac{T_2}{T_1} = \frac{22.4 \times 373}{273} = 30.6L$$

8. (a) $V_{mss} = \sqrt{\frac{3RT}{M}}$; $V_{mss} = \sqrt{T}$
Given, $V_1 = V$, $T_1 = 300K$
 $V_2 = 2V$, $T_2 = ?$
 $= \frac{V_1}{V_2} = \sqrt{\frac{T_1}{T_2}} = \left(\frac{V}{2V}\right)^2 = \frac{300}{T_2}$
 $T_2 = 300 \times 4 = 1200K$

9. (d) When two vessels are joined together, the volume will be doubled hence effective pressure will be halved

$$P = \frac{P_1 + P_2}{2} = \frac{100 + 400}{2} = 250mm$$

10. (a) $P_1V_1 = P_2V_2$ at constant *T*

$$10^4 \cdot 100 = 10^5 \times V_2$$

$$V_2 = 10cc$$

 (b) Kinetic energy will also remain constant if Temperature is constant.

12. (a)
$$\frac{r_{SO_2}}{r_{O_2}} = \sqrt{\frac{M_{O_2}}{M_{SO_2}}} = \sqrt{\frac{32}{64}} = \frac{1}{\sqrt{2}}$$

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