

**Objective Questions** 

# Characteristics and Measurable properties of gases

- **1.** Which one of the following statements is not correct about the three states of matter *i.e.* solid, liquid and gaseous
  - (a) Molecules of a solid possess least energy whereas those of a gas possess highest energy
  - (b) The density of solid is highest whereas that of gases is lowest
  - (c) Gases like liquids possess definite volumes
  - (d) Molecules of a solid possess vibratory motion
- **2.** The temperature and pressure at which ice, liquid water and water vapour can exist together are
  - (a)  $0^{\circ} C$ , 1 atm
- (b)  $2^{\circ} C$ , 4.7 atm
- (c)  $0^{\circ} C$ , 4.7 mm
- (d)  $-2^{\circ}C$ , 4.7 mm
- 3. Which of the following is true about gaseous state
  - (a) Thermal energy = Molecular attraction
  - (b) Thermal energy >> Molecular attraction
  - (c) Thermal energy << Molecular attraction
  - (d) Molecular forces >> Those in liquids
- 4. Kinetic energy of molecules is highest in
  - (a) Gases
- (b) Solids
- (c) Liquids
- (d) Solutions
- **5.** Which of the following statement is correct
  - (a) In all the three states the molecules possess random translational motion
  - (b) Gases cannot be converted into solids without passing through liquid state
  - (c) One of the common property of liquids and gases is viscosity
- (d) According to Boyle's law V/P is constant at constant T
- **6.** A volume of 1  $m^3$  is equal to
  - (a)  $1000 cm^3$
- (b)  $100 cm^3$
- (c)  $10 dm^3$
- (d)  $10^6 cm^3$
- **7.** Which one of the following is not a unit of pressure
  - (a) Newton
- (b) Torr
- (c) Pascal
- (d) Bar
- **8.**  $1^{\circ}C$  rise in temperature is equal to a rise of
  - (a)  $1^{o} F$

- (b)  $9/5^{\circ}F$
- (c)  $5/9^{o}F$
- (d)  $33^{o}F$

- **9.** Which of the following relations for expressing volume of a sample is not correct
  - (a)  $1L = 10^3 \, ml$
- (b)  $1 dm^3 = 1 L$
- (c)  $1L = 10^3 m^3$
- (d)  $1L = 10^3 cm^3$
- **10.** One atmosphere is numerically equal to approximately
  - (a)  $10^6 \text{ dynes } cm^{-2}$
- (b)  $10^2 \text{ dynes } cm^{-2}$ (d)  $10^8 \text{ dynes } cm^{-2}$
- (c)  $10^4 \text{ dynes } cm^{-2}$
- -
- **11.** 2gm of  $O_2$  at  $27^{\circ}C$  and 760mm of Hg pressure has volume [BCECE 2005]
  - (a) 1.5 lit.
- (b) 2.8 lit.
- (c) 11.2 lit.
- (d) 22.4 lit.
- 12. Pressure of a gas in a vessel can be measured by
  - (a) Barometer
- (b) Manometer
- (c) Stalgometer
- (d) All the baove
- 13. Volume occupied by a gas at one atmospheric pressure and  $0^{o}C$  is V mL. Its volume at 273 K will be

#### [Bihar MADT 1982]

- (a) *V ml*
- (b) V/2 ml

(c) 2 V

- (d) None of these
- **14.** Which one of the following statements is wrong for gases

#### [CBSE PMT 1999]

- (a) Gases do not have a definite shape and volume
  - (b) Volume of the gas is equal to the volume of the container confining the gas
  - (c) Confined gas exerts uniform pressure on the walls of its container in all directions
  - (d) Mass of the gas cannot be determined by weighing a container in which it is enclosed
- **15.** Which of the following exhibits the weakest intermolecular forces [AIIMS 2000]
  - (a)  $NH_3$
- (b) *HCl*

(c) *He* 

- (d)  $H_2O$
- 16.  $N_2$  is found in a litre flask under  $100\,kPa$  pressure and  $O_2$  is found in another 3 litre flask under  $320\,kPa$  pressure. If the two flasks are connected, the resultant pressures is

#### [Kerala PMT 2004]

- (a) 310 *kPa*
- (b) 210 kPa
- (c) 420 kPa
- (d) 365 kPa
- (e) 265 kPa

# Ideal gas equation and Related gas laws

**1.** If *P*, *V*, *T* represent pressure, volume and temperature of the gas, the correct representation of Boyle's law is

[BIT Ranchi 1988]

- (a)  $V \propto \frac{1}{T}$  (at constant *P*)
- PV = RT
- (c)  $V \propto 1/P$  (at constant T) (d)
- PV = nRT
- **2.** At constant temperature, in a given mass of an ideal gas

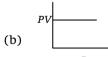
#### [CBSE PMT 1991]

- (a) The ratio of pressure and volume always remains constant
- (b) Volume always remains constant
- (c) Pressure always remains constant
- (d) The product of pressure and volume always remains constant
- **3.** Air at sea level is dense. This is a practical application of

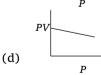
#### [Kerala CEE 2000]

- (a) Boyle's law
- (b) Charle's law
- (c) Avogadro's law
- (d) Dalton's law
- **4.** If  $20 cm^3$  gas at 1 atm. is expanded to 50  $cm^3$  at constant T, then what is the final pressure[CPMT 1988]
  - (a)  $20 \times \frac{1}{50}$
- (b)  $50 \times \frac{1}{20}$
- (c)  $1 \times \frac{1}{20} \times 50$
- (d) None of these
- 5. Which of the following statement is false[BHU 1994]
  - (a) The product of pressure and volume of fixed amount of a gas is independent of temperature
  - (b) Molecules of different gases have the same K.E. at a given temperature
  - (c) The gas equation is not valid at high pressure and low temperature
  - (d) The gas constant per molecule is known as Boltzmann constant
- **6.** Which of the following graphs represent Boyle's law









- 7. Densities of two gases are in the ratio 1 : 2 and their temperatures are in the ratio 2 : 1, then the ratio of their respective pressures is [BHU 2000]
  - (a) 1:1
- (b) 1:2
- (c) 2:1
- (d) 4:1
- **8.** At constant pressure, the volume of fixed mass of an ideal gas is directly proportional to [EAMCET 1985]
  - (a) Absolute temperature
- (b) Degree centigrade

- (c) Degree Fahrenheit (d) None
- **9.** Which of the following expression at constant pressure represents Charle's law [AFMC 1990]
  - (a)  $V \propto \frac{1}{T}$
- (b)  $V \propto \frac{1}{T^2}$
- (c)  $V \propto T$
- (d)  $V \propto d$
- 10. Use of hot air balloons in sports and meteorological obsevations is an application of [Kerala ME
  - (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  $273^{\,o}C$  to  $0^{\,o}C$  keeping the volume constant, its pressure would become
  - (a) 1/2 atm
- (b) 1/273 atm
- (c) 2 atm
- (d) 273 atm
- Pressure remaining the same, the volume of a given mass of an ideal gas increases for every degree centigrade rise in temperature by definite fraction of its volume at

#### [CBSE PMT 1989]

- (a) 0°C
- (b) Its
- critical

temperature

- (c) Absolute zero temperature
- (d) Its
- Boyle
- 13. A certain sample of gas has a volume of 0.2 *litre* measured at 1 *atm*. pressure and  $0^{\circ}C$ . At the same pressure but at  $273^{\circ}C$ , its volume will be **EAMCET 1992, 9** 
  - (a) 0.4 litres
- (b) 0.8 litres
- (c) 27.8 litres
- (d) 55.6 litres
- **14.** 400  $cm^3$  of oxygen at  $27^{\circ}C$  were cooled to  $-3^{\circ}C$  without change in pressure. The contraction in volume will be
  - (a)  $40 cm^3$
- (b)  $30 \text{ cm}^3$
- (c)  $44.4 \text{ cm}^3$
- (d)  $360 \text{ cm}^3$
- **15.** The pressure p of a gas is plotted against its absolute temperature T 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

**34.** At  $0^{o}C$  and one atm pressure, a gas occupies 100

gas will be

cc. If the pressure is increased to one and a half-

time and temperature is increased by one-third of

absolute temperature, then final volume of the

			Ga	aseous State 237
	$(2)$ $(2P_1T_1)$ $(2P_1T_1)$		(a) 1.987 cal K <sup>-1</sup> mol <sup>-1</sup>	(b) 8.3 cal K <sup>-1</sup> mol <sup>-1</sup>
	(a) $\frac{2P_1T_1}{T_1+T_2}$ (b) $\frac{T_1}{2P_1T_2}$		(c) $0.0821 \ lit \ K^{-1} \ mol^{-1}$	(d) 1.987 Joules $K^{-1}$ $mol^{-1}$
	2P.T <sub>2</sub> 2P.	25.	The constant $R$ is	[Orissa 1990]
	(c) $\frac{2P_1T_2}{T_1+T_2}$ (d) $\frac{2P_1}{T_1+T_2}$		(a) Work done per mol	ecule
10	"One gram molecule of a gas at N.T.P. occupies		(b) Work done per deg	ree absolute
17.	22.4 <i>litres.</i> " This fact was derived from[CPMT 1981, 19	00=1	(c) Work done per deg	ree per mole
	(a) Dalton's theory	וכפפ	(d) Work done per mol	_
	(b) Avogadro's hypothesis	26.	-	ement. In the gas equation,
			PV = nRT	[CBSE PMT 1992]
	(c) Berzelius hypothesis		(a) <i>n</i> is the number of	molecules of a gas
- 0	(d) Law of gaseous volume		(b) V denotes volume of	of one mole of the gas
18.	In a closed flask of 5 <i>litres</i> , 1.0 $g$ of $H_2$ is heated		(c) <i>n</i> moles of the gas l	have a volume V
	from 300 to 600 <i>K</i> . which statement is not correct[ <b>c</b>	BSE PN	IT(d991) is the pressure	of the gas when only one
	(a) Pressure of the gas increases		mole of gas is prese	ent
	(b) The rate of collision increases	27.	The correct value of the	e gas constant R is close to
	(c) The number of moles of gas increases			[CBSE PMT 1992]
	(d) The energy of gaseous molecules increases		(a) 0.082 litre-atmops	here K
19.	Which one of the following statements is false		(b) 0.082 litre-atmospi	here $K^{-1} mol^{-1}$
	[Manipal PMT 1991]		-	
	(a) Avogadro number = $6.02 \times 10^{21}$		(c) 0.082 litre-atmosphe	ere -1 K mole -1
	(b) The relationship between average velocity $(\overline{v})$		(d) $0.082$ litre <sup>-1</sup> atmosph	ere <sup>–1</sup> K mol
	and root mean square velocity $(u)$ is	20	_	
	$\overline{v} = 0.9213 \ u$	28.	0	
	(c) The mean kinetic energy of an ideal gas is		(a) $0.0821$ litre atm $K^{-1}$	
	independent of the pressure of the gas		(b) 2 calories K <sup>-1</sup> mole	
	(d) The root mean square velocity of the gas can		(c) 8.31 <i>joule K</i> <sup>-1</sup> <i>mole</i>	1
	be calculated by the formula $(3RT/M)^{1/2}$		(d) None	
	•	29.	Gas equation $PV = nRT$	is obeyed by [BHU 2000]
20.	The compressibility of a gas is less than unity at	_	-	ocess (b)Only adiabatic proces
	STP. Therefore [IIT 2000]		(c) Both (a) and (b)	(d) None of these
	(a) $V_m > 22.4 \ litres$ (b) $V_m < 22.4 \ litres$	30.		nber of moles per litre in
	(c) $V_m = 22.4 \ litres$ (d) $V_m = 44.8 \ litres$	30.	_	e P, gas constant R and
21.	In the equation of sate of an ideal gas $PV = nRT$ ,		temperature <i>T</i> is	e 1, gas constant it and
	the value of the universal gas constant would		1	[AIEEE 2002]
	depend only on		(a) <i>PT/R</i>	(b) PRT
	[KCET 2005]		(c) <i>P/RT</i>	(d) <i>RT/P</i>
	(a) The nature of the gas	31.		eal gas at 546 K occupy a
	(b) The pressure of the gas		volume of 44.8 litres, t	_
	(c) The units of the measurement			[NCERT 1981; JIPMER 1991]
	(d) None of these		(a) 2 atm	(b) 3 atm
22.	In the ideal gas equation, the gas constant $R$ has		(c) 4 atm	(d) 1 atm
	the dimensions of [NCERT 1982]	32.		Ie gas occupy 22.4 litres at
	(a) $mole-atm K^{-1}$ (b) $litre mole$		-	neric pressure [KCET 1992]
	(c) litre-atm K <sup>-1</sup> mole <sup>-1</sup> (d) erg K <sup>-1</sup>		(a) 0.90	(b) 1.11
23.	In the equation $PV = nRT$ , which one cannot be		(c) 0.11	(d) 1.0
	the numerical value of <i>R</i> [BIT 1987]	33.		a gas at 1 <i>atm</i> . pressure and
	(a) $8.31 \times 10^7 erg \ K^{-1} mol^{-1}$		273 K is	F=
	(b) $8.31 \times 10^7$ dyne cm $K^{-1}$ mol <sup>-1</sup>		(a) an A litron	[EAMCET 1992]
	·		(a) 22.4 litres (c) 44.8 litres	(b) 11.2 litres (d) 5.6 litres
	(c) $8.31 \ JK^{-1} mol^{-1}$		(6) 44.0 111163	(4) 5.0 111103

(d)  $8.31 \text{ atm. } K^{-1} \text{mol}^{-1}$ 

the gas constant R

24.

Which one of the following indicates the value of

[EAMCET 1989]

following temperatures will its density become

A sample of gas occupies 100 ml at  $27^{\circ}C$  and 740

mm pressure. When its volume is changed to 80

(b)  $30^{\circ}C$ 

(d) 300 K

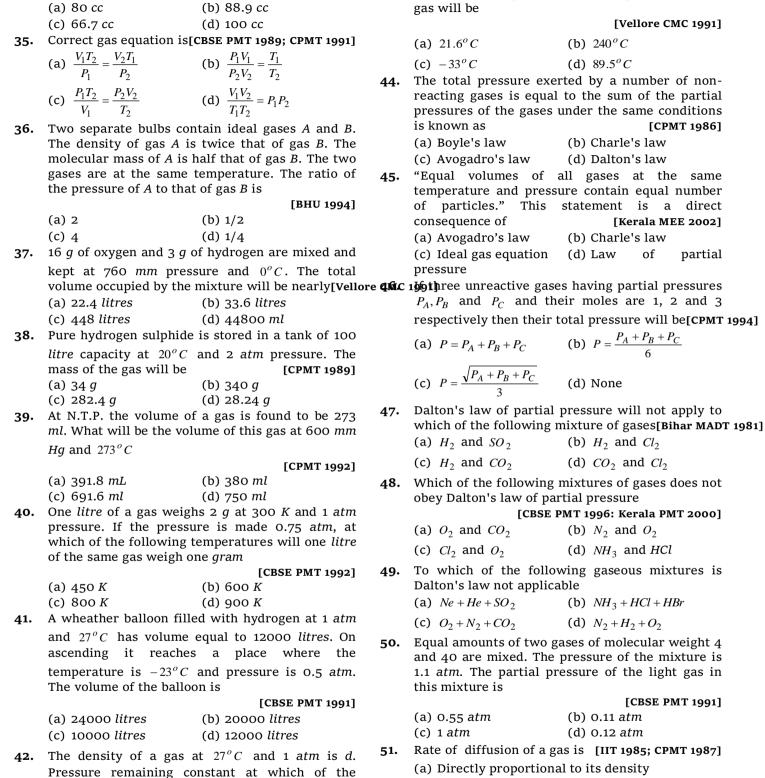
[CBSE PMT 1992]

0.75 d

43.

(a)  $20^{\circ} C$ 

(c) 400 K



[DCE 2000]

ml at 740 mm pressure, the temperature of the

(b) Directly proportional to its molecular mass

molecular mass

molecular mass

of diffusion

(c) Directly proportional to the square root of its

(d) Inversely proportional to the square root of its

Which of the following gas will have highest rate

[IIT 1996]

	[Pb. CET Sa	ample paper 1993; CPMT 1990]	62.	At constant volume and temperature conditions,
	(a) $NH_3$	(b) $N_2$		the rate of diffusion $D_A$ and $D_B$ of gases $A$ and $B$
	(c) <i>CO</i> <sub>2</sub>	(d) $O_2$		having densities $ ho_A$ and $ ho_B$ are related by the
53.	Which of the follows	ing relationship is correct,		expression [IIT 1993]
	its density	diffusion of a gas and $d$ is [CPMT 1994]		(a) $D_A = \left[D_B \cdot \frac{\rho_A}{\rho_B}\right]^{1/2}$ (b) $D_A = \left[D_B \cdot \frac{\rho_B}{\rho_A}\right]^{1/2}$
	(a) $r \propto \sqrt{1/d}$	(b) $r \propto \sqrt{d}$		$(a)^{1/2}$ $(a)^{1/2}$
	(c) $r=d$	(d) $r \propto d$		(c) $D_A = D_B \left(\frac{\rho_A}{\rho_B}\right)^{1/2}$ (d) $D_A = D_B \left(\frac{\rho_B}{\rho_A}\right)^{1/2}$
54.		man's law at a given	63.	Atmolysis is a process of
	$r_A / r_B$ of gases A and E	o of the rates of diffusion  is given by [IIT 1998]	-3.	(a) Atomising gas molecules
				(b) The breaking of atoms to sub-atomic particles
	(a) $(P_A / P_B)(M_A / M_B)^{1/2}$	_		(c) Separation of gases from their gaseous
	(b) $(M_A / M_B)(P_A / P_B)^{1/2}$	2		mixture
	(c) $(P_A / P_B)(M_B / M_A)^{1/2}$	2	64.	(d) Changing of liquids to their vapour state  A bottle of ammonia and a bottle of dry hydrogen
			04.	chloride connected through a long tube are
	(d) $(M_A / M_B)(P_B / P_A)^{1/2}$			opened simultaneously at both ends, the white
		the pressures and molecular		ammonium chloride ring first formed will be[IIT 1988]
	weights of gases A and	• •		(a) At the centre of the tube
55.		te of diffusion of a given elium is 1.4. The molecular		(b) Near the hydrogen chloride bottle
	weight of the element			(c) Near the ammonia bottle
		[Kerala PMT 1990]	65.	(d) Throughout the length of the tube Which of the following pairs will diffuse at the
	(a) 2	(b) 4	05.	same rate through a porous plug [EAMCET 1990]
	(c) 8	(d) 16		(a) $CO, NO_2$ (b) $NO_2, CO_2$
56.	A gas diffuse 1/5 time molecular weight is	les as fast as hydrogen. Its [CPMT 1992; Bihar CEE 1982]		(c) $NH_3, PH_3$ (d) $NOC_2H_6$
	(a) 50	(b) 25	66.	If 4 $g$ of oxygen diffuse through a very narrow
	<u></u>	<u>_</u>		hole, how much hydrogen would have diffused
	(c) $25\sqrt{2}$	(d) $50\sqrt{2}$		under identical conditions [CPMT 1971] (a) $16 g$ (b) $1 g$
57•		at of a gas which diffuses ag at 1/6th of the speed of		(a) 16 <i>g</i> (b) 1 <i>g</i> (c) 1/4 <i>g</i> (d) 64 <i>g</i>
		cal conditions is[EAMCET 1990]	67.	A gas diffuse at a rate which is twice that of
	(a) 27	(b) 72	٠,٠	another gas $B$ . The ratio of molecular weights of $A$
	(c) 36	(d) 48		to <i>B</i> is <b>[EAMCET 1986]</b>
58.	Molecular weight of a	gas that diffuses twice as		(a) 1.0 (b) 0.75
	rapidly as the gas with	n molecular weight 64 is[EAMCE	ET 199	4(c) 0.50 (d) 0.25
	(a) 16	(b) 8	68.	Two grams of hydrogen diffuse from a container in 10 <i>minutes</i> . How many <i>grams</i> of oxygen would
	(c) 64	(d) 6.4		diffuse through the same container in the same
59.		rogen and oxygen are 0.09		time under similar conditions [MNR 1980]
	and 1.44 $g$ $L^{-1}$ . If the	rate of diffusion of hydrogen		(a) $0.5 g$ (b) $4 g$
		n in the same units will be[RPM		
	(a) 4 (c) 16	(b) 1/4 (d) 1/16	69.	The rate of diffusion of methane at a given
60.		A is 5 times that of B, what		temperature is twice that of $X$ . The molecular weight of $X$ is
	will be the density rat			[MNR 1995; Kerala CEE 2001]
	(a) 1/25	(b) 1/5		(a) 64.0 (b) 32.0
	(c) 25	(d) 4		(c) 40.0 (d) 80.0
61.		gases are in the ratio of 1:	70.	$X \ ml$ of $H_2$ gas effuses through a hole in a
		ates of diffusion is[CPMT 1995]		container in 5 seconds. The time taken for the
	(a) 16:1	(b) 4:1		effusion of the same volume of the gas specified
	(c) 1:4	(d) 1:16		below under identical condition is

	240 Gaseous sta	te				
	(a) 10 seconds : He	(b) 20 seconds : O <sub>2</sub>		(b) $\frac{1}{2}$ that of $SO_2$		
	(c) 25 seconds : CO	(d) 55 seconds : $CO_2$		<u> </u>		
71.	At what temperature,	the rate of effusion of $N_2$		(c) Twice that of $SO_2$		
	would be 1.625 times the	hat of $SO_2$ at $50^{o}C$ [CBSE PMT:	1996]	(d) One fourth that of So	$O_2$	
	(a) 110 <i>K</i>	(b) 173 <i>K</i>	81.	Five grams each of the	following gases at $87^{\circ}C$	
	(c) 373 K	(d) 273 <i>K</i>			are taken. Which of them	
72.		$C(s) + H_2O(l) \rightarrow CO(g) + H_2(g)$		will have the least volur	me [MNR 1991] (b) <i>HCl</i>	
	from 48.0 <i>g</i> of carbon	f the gases produced at STP		(a) HF (c) HBr	(d) HI	
	(a) 179.2 <i>L</i>	(b) 89.6 L	82.		ng scientists has not done	
	(c) 44.8 <i>L</i>	(d) 22.4 <i>L</i>			gases [Bihar MADT 1980]	
73.		ccupies a volume of 2.24 $L$ ,		(a) Boyle	(b) Charles	
	the gas can be	[Haryana CEET 2000]		(c) Avogadro	(d) Faraday	
	(a) $O_2$	(b) <i>CO</i>	83.		equal number of nitrogen	
	(c) $NO_2$	(d) <i>CO</i> <sub>2</sub>			at a pressure of <i>P mm</i> . If rom the system then the	
74.		s will a pure sample of an		pressure will be	tom the system then the	
	ideal gas not only exhi	bit a pressure of 1 atm but			[MP PMT 1985]	
				(a) <i>P</i>	(b) 2 <i>P</i>	
	$(R = 0.082 \text{ litre atm mol}^{-1} \text{ d})$	$eg^{-1}$ ) [CBSE PMT 1993]		(c) P/2	(d) $P^2$	
	(a) At STP (b) When $V = 22.4$ litres		84.		car are filled to the same and <i>Ne</i> separately, then	
	(c) When $T = 12 K$			which one will be filled	first [Manipal PMT 2001]	
	(d) Impossible under a	ny conditions		(a) $N_2$	(b) $O_2$	
75.	There are $6.02 \times 10^{22}$ m	olecules each of $N_2, O_2$ and		(c) $H_2$	(d) Ne	
	$H_2$ which are mixed to	ogether at 760 mm and 273	85.	Which of the following	ing gas mixture is not	
		ture in grams is [Pb. PMT 1997]		= =	aw of partial pressure[Pb. CE	
	(a) 6.2	(b) 4.12		(a) $SO_2$ and $Cl_2$	(b) $CO_2$ and $N_2$	
76.	(c) 3.09 Volume of 4.4 <i>g</i> of <i>CO</i> <sub>2</sub>	(d) 7 at NTP is <b>[Pb. CET 1997</b> ]		(c) $CO$ and $CO_2$	(d) $CO$ and $N_2$	
, <b>0.</b>	(a) 22.4 $L$	(b) 44.8 <i>L</i>	86.		intity of gas will occupy a	
	(a) 22.4 L (c) 2.24 L	(d) 4.48 <i>L</i>			occupies a volume of 100ml	
77.	The energy of an ideal			constant):	n? (while temperature is [Pb. CET 2000]	
	(a) Pressure	(b) Volume		(a) 700 mm	(b) 800 mm	
	(c) Number of moles	(d) Temperature		(c) 100 mm	(d) 1200 mm	
78.		contains 200 $ml$ liquid in lar. Suppose $CO_2$ behaves	87.	At constant temperatur	e and pressure which gas	
	like an ideal gas, the vo	olume of the dissolved $CO_2$		will diffuse first $H_2$ or	O <sub>2</sub> ? [Pb. CET 2000]	
	at STP is	[CBSE PMT 1991]		(a) Hydrogen		
	(a) 0.224 litre	(b) 0.448 litre		<ul><li>(b) Oxygen</li><li>(c) Both will diffuse in s</li></ul>	same time	
	(c) 22.4 litre	(d) 2.24 litre		(d) None of the above	Junic tillic	
79.		a gas is 11.2. The volume is gas at N.T.P. is	88.	When a jar containing gaseous mixture of equal		
	occupied by 11.2 g of th	=		volumes of $CO$ and $U$	is placed in a colution of	
		[MNR 1982; CBSE PMT 1991]			is placed in a solution of olution level will Ph. CET 200	
	(a) 1 <i>L</i> (c) 22.4 <i>L</i>	=			is placed in a solution of olution level will[Pb. CET 200 (b) Fall	

N.T.P. and weighted. It was then evacuated, filled with  $SO_2$  at the same temperature and pressure,

(a) The same as that of  $SO_2$ 

and again weighted. The weight of oxygen will be [NCERT 1989]) 22.4 litre

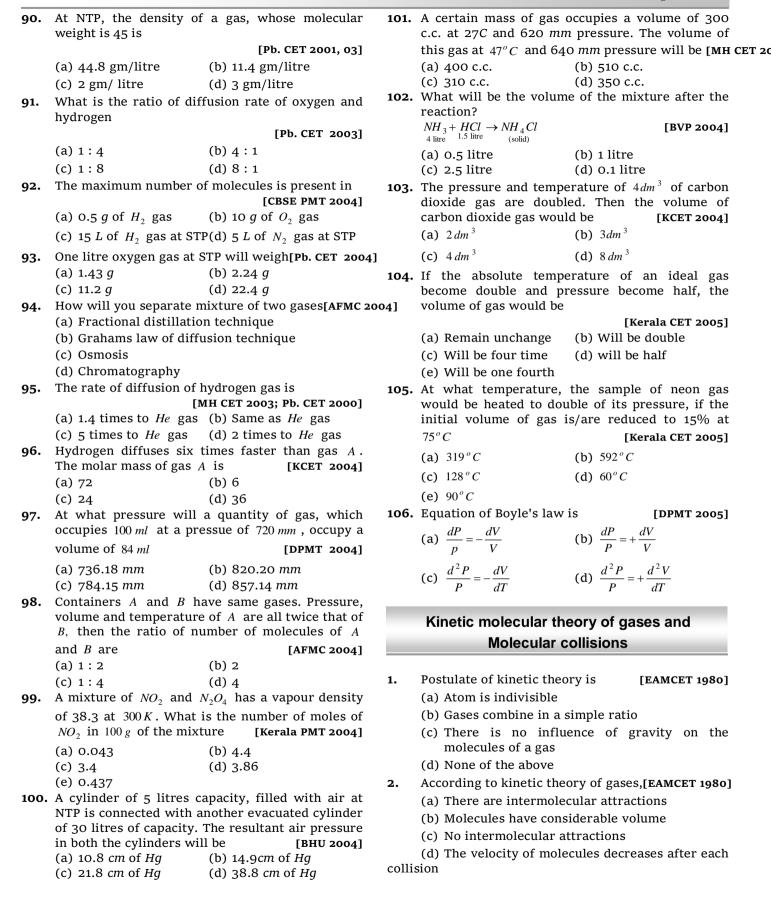
**89.** At S.T.P.  $1g CaCO_3$  on decomposition gives  $CO_2$ 

(c) 0.224 litre

[Pb. CET 2000]

(b) 2.24 litre

(d) 11.2 litre



In deriving the kinetic gas equation, use is made 3. of the root mean square velocity of the molecules because it is

#### [Bihar MADT 1980]

- (a) The average velocity of the molecules
- (b) The most probable velocity of the molecules
- (c) The square root of the average square velocity of the molecules
- (d) The most accurate form in which velocity can be used in these calculations
- Kinetic energy of a gas depends upon its[Bihar 4. MADT 1982]
  - (a) Molecular mass
- (b) Atomic mass
- (c) Equivalent mass
- (d) None of these
- The kinetic theory of gases perdicts that total 5. kinetic energy of a gaseous assembly depends on [NCERT 1984] Have equal average kinetic energies
  - (a) Pressure of the gas
  - (b) Temperature of the gas
  - (c) Volume of the gas
  - (d) Pressure, volume and temperature of the gas
- 6. According to kinetic theory of gases, the energy per mole of a gas is equal to [EAMCET 1985]
  - (a) 1.5 RT
- (b) RT
- (c) 0.5 RT
- (d) 2.5 RT
- 7. Internal energy and pressure of a gas per unit volume are related as [CBSE PMT 1993]
  - (a)  $P = \frac{2}{3}E$
- (b)  $P = \frac{3}{2}E$
- (c)  $P = \frac{1}{2}E$
- (d) P = 2E
- 8. The translational kinetic energy of an ideal gas depends only on its
  - (a) Pressure
- (b) Force
- (c) Temperature
- (d) Molar mass
- Helium atom is two times heavier than a 9. hydrogen molecule at 298 K, the average kinetic energy of helium is [IIT 1982]
  - (a) Two times that of a hydrogen molecule
  - (b) Same as that of a hydrogen molecule
  - (c) Four times that of a hydrogen molecule
  - (d) Half that of a hydrogen molecule
- Which of the following is valid at absolute zero

#### [Pb. CET 1985]

- (a) Kinetic energy of the gas becomes zero but the molecular motion does not become zero
- (b) Kinetic energy of the gas becomes zero and molecular motion also becomes zero
- (c) Kinetic energy of the gas decreases but does not become zero
- (d) None of the above
- The average K.E. of an ideal gas in calories per 11. mole is approximately equal to [EAMCET 1989]
  - (a) Three times the absolute temperature
  - (b) Absolute temperature
  - (c) Two times the absolute temperature
  - (d) 1.5 times the absolute temperature

12. According to kinetic theory of gases, for a diatomic molecule

[MNR 1991]

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

[CBSE PMT 1993, 2000]

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

[CBSE PMT 2000]

- (a)  $\overline{KE}_{CO} = \overline{KE}_{N}$ ,
- (b)  $\overline{KE}_{CO} > \overline{KE}_{N_2}$
- (c)  $\overline{KE}_{CO} < \overline{KE}_{N_2}$
- (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  $CO_2$
  - (d) The total translational  $\mathit{KE}$  of both  $\mathit{N}_2$  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 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

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

Which of the following has maximum root mean square velocity at the same temperature[Manipal PMT 200

(b) *CO*<sub>2</sub>

(d)  $H_2$ 

(d)  $\sqrt{2}:\sqrt{8/\pi}:\sqrt{3}$ 

(2) A negative value of  $\mu_{JT}$  (Joule Thomson

coefficient corresponds to warming of a gas

	then for an ideal gas, th	ie density is given by <b>[cbse pm</b>	T 1989	9, 91] on expansion.
	(a) $\frac{RT}{PM}$	(b) $\frac{P}{RT}$		(3) The temperature at which neither cooling nor
				heating effect is observed is known as
	(c) $\frac{M}{V}$	(d) $\frac{PM}{RT}$		inversion temperature.
	•	KI		Which of the above statements are correct
19.	_	naximum density when[ <b>cpmT</b> :	2000]	(a) 1 and 2 (b) 1 and 3
	(a) $P = 0.5 atm, T = 600 K$		-0	(c) 2 and 3 (d) 1, 2 and 3
	(b) $P = 2 atm, T = 150 K$		28.	Vibrational energy is [Pb. CET 1985]
	(c) $P = 1 atm, T = 300 K$			(a) Partially potential and partially kinetic
	(d) $P = 1.0 atm, T = 500 K$			(b) Only potential
	,			(c) Only kinetic (d) None of the above
20.		rature of a gas is $-80^{\circ}C$ ,	20	
	effect at	oling under Joule-Thomson	29.	At the same temperature and pressure, which of the following gases will have the highest kinetic
	(a) 298 <i>K</i>	(b) 273 <i>K</i>		energy per mole
	(c) 193 K	(d) 173 <i>K</i>		[MNR 1991]
21.		a gas $X'$ is 1.4. The number		(a) Hydrogen (b) Oxygen
	•	present in 11.2 <i>litres</i> of it at		(c) Methane (d) All the same
	N.T.P. is	or coord in 11.2 tier of or it de	30.	Dimensions of pressure are the same as that of
		[CBSE PMT 1989]		[CBSE PMT 1995]
	(a) $6.02 \times 10^{23}$	(b) $1.2 \times 10^{24}$		(a) Energy (b) Force
	(c) $3.01 \times 10^{23}$	(d) $2.01 \times 10^{23}$		(c) Energy per unit volume (d)Force per unit volume
22.	` '	0.00130 $q/ml$ . The vapour	31.	The density of a gas An is three times that of a gas
	density of air will be	[DCE 2000]		B. if the molecular mass of A is $M$ , the molecular
	(a) 0.00065	(b) 0.65		mass of $B$ is [CPMT 1987]
	(c) 14.4816	(d) 14.56		
23.		the density of liquid water		(a) 3 $M$ (b) $\sqrt{3} M$
23.		f water vapour is 0.0006 g		(c) $M/3$ (d) $M/\sqrt{3}$
	•	-		Malagular angodo
		me occupied by water steam at that temperature		Molecular speeds
	is	[IIT 2000]	1.	The ratio of root mean square velocity to average
	(a) 6 $cm^3$	(b) 60 cm <sup>3</sup>		velocity of gas molecules at a particular
				temperature is [IIT 1981]
	(c) $0.6 \text{ cm}^3$	(d) 0.06 cm <sup>3</sup>		(a) 1.086:1 (b) 1:1.086
24.	The ratio $\gamma$ for inert gas			(c) 2:1.086 (d) 1.086:2
	(a) 1.33	(b) 1.66	2.	Which is not true in case of an ideal gas[CBSE PMT 1991]
	(c) 2.13	(d) 1.99	_	(a) It cannot be converted into a liquid
25.	The density of neon wil	l be highest at[CBSE PMT 1990]	J	(b) There is no interaction between the molecules
	(a) S.T.P.	(b) $0^{\circ} C, 2 atm$		(c) All molecules of the gas move with same
	(c) 273°C,1 atm	(d) 273°C, 2 atm	spee	<u> </u>
26	,	, ,		(d) At a given temperature, PV is proportional to
26.	Absolute zero is defined	-		the amount of the gas
	(a) At which all molecu	[CBSE PMT 1990]	3.	The ratio among most probable velocity, mean
	(a) At which an molecu	iai inotion ccases		velocity and root mean square velocity is given by [CBSE

(a) 1:2:3

(a)  $SO_2$ 

(c)  $O_2$ 

(c)  $\sqrt{2}:\sqrt{3}:\sqrt{8/\pi}$ 

**18.** If P, V, M, T and R are pressure, volume, molar mass, temperature and gas constant respectively,

(b) At which liquid helium boils

**27.** Consider the following statements :

(1) Joule-Thomson experiment is isoenthalpic as

(c) At which ether boils

well as adiabatic.

(d) All of the above

(c) Increase in molecular attraction(d) Decrease in mean free path

- What is the pressure of 2 mole of  $NH_3$  at  $27^{\circ}C$ 24. when its 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
- The root mean square velocity of one mole of a 25. monoatomic having molar mass M is  $U_{ms}$ . The relation between the average kinetic energy (E) of the  $U_{rms}$  is

[IIT-JEE Screening 2004]

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

(b) 
$$U_{rms} = \sqrt{\frac{2E}{3M}}$$

(c) 
$$U_{rms} = \sqrt{\frac{2E}{M}}$$

(d) 
$$U_{rms} = \sqrt{\frac{E}{3M}}$$

26. Ratio of average to most probable velocity is

[Orissa JEE 2005]

- (a) 1.128
- (b) 1.224
- (c) 1.0
- (d) 1.112
- If the  $v_{ms}$  is  $30R^{1/2}$  at  $27^{\circ}C$  then calculate the molar mass of gas in kilogram. [DPMT 2005]
  - (a) 1

(b) 2

(c) 4

(d) 0.001

# Real gases and Vander waal's equation

equation explains The Vander Waal's 1. behaviour of

[DPMT 1981]

- (a) Ideal gases
- (b) Real gases
- (c) Vapour
- (d) Non-real gases
- 2. Gases deviate from the ideal gas behaviour because their molecules [NCERT 1981]
  - (a) Possess negligible volume
  - (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 3. Z = PV / RT. The compressibility factor of ideal gas is

[Pb. CET 1986]

(a) o

(b) Infinity

(c) 1

- (d) -1
- In Vander Waal's equation of state for a non-ideal

gas, the term that 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 5. gases. For n moles 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) 
$$\left(P + \frac{n^2 a}{V^2}\right)(V - nb) = nRT$$

Any gas shows maximum deviation from ideal gas

[CPMT 1991]

- (a)  $0^{\circ}C$  and 1 atmospheric pressure
- (b)  $100^{\circ}C$  and 2 atmospheric pressure
- (c)  $-100^{\circ}C$  and 5 atmospheric pressure
- (d)  $500^{\circ}C$  and 1 atmospheric pressure
- The temperature at which the second virial 7. coefficient of real gas is zero is called [AFMC 1993]
  - (a) Critical temperature
  - (b) Eutetic point
  - (c) Boiling point
  - (d) Boyle's temperature
- When is deviation more in the behaviour of a gas 8. from the ideal gas equation PV = nRT

#### [DPMT 1981; NCERT 1982; CBSE PMT 1993]

- (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
  - (c) Shape and repulsive forces of molecules
  - (d) Attractive force and volume of the molecules
- Gas deviates from ideal gas nature because molecules

[CPMT 1996]

- (a) Are colourless
- (b) Attract each other
- (c) Contain covalent bond
- (d) Show Brownian movement
- The Vander Waal's equation reduces itself to the 11. ideal gas equation at [Kerala MEE 2001; CBSE PMT 2002]
  - (a) High pressure and low temperature
  - (b) Low pressure and low temperature
  - (c) Low pressure and high temperature
  - (d) High pressure and high temperature
- The compressibility factor for an ideal gas is[IIT 1997] 12.
  - (a) 1.5
- (b) 1.0
- (c) 2.0
- (d) ∞
- When an ideal gas undergoes unrestrained 13. 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 14. 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
- (c) When both the temperature and pressure are high
- (d) When the temperature is high and pressure is low
- A real gas most closely approaches the behaviour 15. of an ideal gas at [KCET 1992]
  - (a) 15 atm and 200 K
- (b) 1 atm and 273 K
- (c) 0.5 atm and 500 K
- (d) 15 atm and 500 K
- 16. The temperature at which real gases obey the ideal gas laws over a wide range of pressure is called

#### [AFMC 1993; IIT 1981, 94]

- (a) Critical temperature
- (b) Boyle temperature
- (c) Inversion temperature
- (d) Reduced temperature
- At low pressure, the Vander Waal's equation is 17.

(a) 
$$Z = \frac{pV_m}{RT} = 1 - \frac{ap}{RT}$$

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

(c) 
$$pV_m = RT$$

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

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, 19. heating effect is observed. Its reason is that [CPMT 1987]
  - (a) Helium is an ideal gas
  - (b) Helium is an inert gas
- (c) The inversion temperature of helium is very low
  - (d) The boiling point of helium is the lowest among the elements

- In van der Waal's equation of state of the gas law, 20. 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
- In which molecule the vander Waal's force is 21. likely to be the most important in determining the m.pt. and b.pt.

[DPMT 2000]

- (a)  $H_2S$
- (b)  $Br_2$
- (c) HCl
- (d) CO
- Pressure exerted by 1 mole of methane in a 0.25 22. litre container at 300K using vander Waal's equation

$$1 = 2.253 \ atm \ l^2 \ mol^{-2}, b = 0.0428 \ lit mol^{-1})$$
 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

- 1. Which set of conditions represents easiest way to liquefy a gas [NCERT 1983]
  - (a) Low temperature and high pressure
  - (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 2. [BHU 1984]
  - (a) Adiabatic expansion of a gas
  - (b) Production of low temperature
  - (c) Production of high temperature
  - (d) None
- An ideal gas can't be liquefied because [CBSE PMT 1992] 3.
  - (a) Its critical temperature is always above  $0^{\circ}C$
  - (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
  - (b) Inversion temperature
  - (c) Critical temperature
  - (d) Room temperature
- An ideal gas obeying kinetic theory of gases can be liquefied if [CBSE PMT 1995]
- (a) Its temperature more than critical is 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 TThe Vander Waal's parameters for gases W, X, Y6. and Z are

Gas	a (atm L² mol <sup>-2</sup> )	b (L mol <sup>-1</sup> )
W	4.0	0.027
X	8.0	0.030
Y	6.0	0.032
Z	12.0	0.027

Which one of these gases has the highest critical temperature

(a) W

(b) X

(c) Y

- (d) Z
- The Vander Waal's constant 'a' for the gases 7.  $O_2, N_2, NH_3$  and  $CH_4$  are 1.3, 1.390, 4.170 and 2.253  $L^2$  atm mol<sup>-2</sup> respectively. The gas which can be most easily liquefied is

[IIT 1989]

- (a)  $O_2$
- (b)  $N_2$
- (c)  $NH_3$
- (d)  $CH_4$
- A gas can be liquefied

[AFMC 2005]

- (a) Above its critical temperature
  - (b) At its critical temperature
  - (c) Below its critical temperature
  - (d) At any temperature
- Which of the following is correct for critical 9. temperature
  - (a) It is the highest temperature at which liquid and vapour can coexist
  - (b) Beyond the critical temperature, there is no distinction between 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
- 10. A gas has a density of 2.68 g/L at stp. Identify the gas
  - (a)  $NO_2$
- (b) Kr
- (c) COS
- (d)  $SO_2$
- Weight of 112 ml of oxygen at NTP on liquefaction 11. would be [DPMT 1984]
  - (a) 0.32 q
- (b) 0.64 q
- (c) 0.16 q
- (d) 0.96 g



- As the temperature is raised from  $20^{\circ} C$  to  $40^{\circ} C$ the average kinetic energy of neon atoms changes by a factor of which of the following [AIEEE 2004]
  - (a) 313/293
- (b)  $\sqrt{(313/293)}$
- (c) 1/2
- (d) 2
- A gas is found to have a formula  $[CO]_r$ . If its 2.. vapour density is 70, the value of x is [DCE 2004]
  - (a) 2.5
- (b) 3.0
- (c) 5.0
- (d) 6.0
- Which of the given sets of temperature and 3. pressure will cause a gas to exhibit the greatest deviation from ideal gas behavior [DCE 2003]
  - (a)  $100^{\circ} C$  and 4 atm
- (b)  $100^{\circ} C$  and 2 atm
- (c)  $-100^{\circ} C$  and 4 atm (d)  $0^{\circ} C$  and 2 atm
- The molecular weight of  $O_2$  and  $SO_2$  are 32 and 64 respectively. If one litre of  $O_2$  at  $15^{\circ}C$  and 750 mm pressure contains 'N' molecules, the number of molecules in two litres of SO2 under the same conditions of temperature and pressure will be [CBSE 1990; MNR 1991]
  - (a) N/2
- (b) N
- (c) 2N
- (d) 4N
- What is the relationship between the average 5. velocity (v), root mean square velocity (u) and most probable velocity (a)

[AFMC 1994]

- (a)  $\alpha: v: u:: 1:1.128:1.224$
- (b)  $\alpha : v : u :: 1.128 : 1 : 1.224$
- (c)  $\alpha: v: u:: 1.128: 1.224: 1$
- (d)  $\alpha : v : u :: 1.124 : 1.228 : 1$
- 6. Consider the following statements: For diatomic gases, the ratio  $C_p / C_v$  is equal to
  - (1) 1.40 (lower temperature)
  - (2) 1.66 (moderate temperature)
  - (3) 1.29 (higher temperature)

which of the above statements are correct

- (a) 1, 2 and 3
- (b) 1 and 2
- (c) 2 and 3
- (d) 1 and 3
- The compressibility factor for an ideal gas is[MP PET 2004 7.
  - (a) 1.5
- (b) 1.0
- (c) 2.0
- (d) ∞
- The compressibility factor of a gas is less than 1 8. at STP. Its molar volume  $V_m$  will be [MP PET 2004]
  - (a)  $V_m > 22.42$
- (b)  $V_m < 22.42$
- (c)  $V_m = 22.42$
- (d) None
- If some moles of  $O_2$  diffuse in 18 sec and same moles of other gas diffuse in 45 sec then what is the molecular weight of the unknown gas[CPMT 1988]

- (a)  $\frac{45^2}{18^2} \times 32$
- (b)  $\frac{18^2}{45^2} \times 32$
- (c)  $\frac{18^2}{45^2 \times 32}$
- (d)  $\frac{45^2}{18^2 \times 32}$
- **10.** The ratio of rates of diffusion of  $SO_2, O_2$  and  $CH_4$  is

[BHU 1992]

- (a)  $1:\sqrt{2}:2$
- (b) 1:2:4
- (c)  $2:\sqrt{2}:1$
- (d)  $1:2:\sqrt{2}$
- 11. If  $C_1, C_2, C_3$ ..... represent the speeds of  $n_1, n_2, n_3$ .... molecules, then the root mean square speed is [IIT 199318.

(a) 
$$\left(\frac{n_1C_1^2 + n_2C_2^2 + n_3C_3^2 + \dots}{n_1 + n_2 + n_3 + \dots}\right)^{1/2}$$

(b) 
$$\frac{(n_1C_1^2 + n_2C_2^2 + n_3C_3^2 + .....)^{1/2}}{n_1 + n_2 + n_3 + .....}$$

(c) 
$$\frac{(n_1C_1^2)^{1/2}}{n_1} + \frac{(n_2C_2^2)^{1/2}}{n_2} + \frac{(n_3C_3^2)^{1/2}}{n_3} + \dots$$

(d) 
$$\left[\frac{(n_1C_1 + n_2C_2 + n_3C_3 + ....)^2}{(n_1 + n_2 + n_3 + ....)}\right]^{1/2}$$

- **12.** 50 *ml* of hydrogen diffuses out through a small hole from a vessel in 20 *minutes*. The time needed for 40 *ml* of oxygen to diffuse out is[CBSE PMT 1994]
  - (a) 12 min
- (b) 64 min
- (c) 8 min
- (d) 32 min
- 13. At what temperature will the average speed of  $CH_4$  molecules have the same value as  ${\cal O}_2$  has at 300 K

[CBSE PMT 1989]

- (a) 1200 K
- (b) 150 K
- (c) 600 K
- (d) 300 K
- 14. A sample of  $O_2$  gas is collected over water at  $23^{\circ}C$  at a barometric pressure of 751 mm Hg (vapour pressure of water at  $23^{\circ}C$  is 21 mm Hg). The partial pressure of  $O_2$  gas in the sample collected is [CBSE PMT 1993]
  - (a) 21 mm Hg
- (b) 751 mm Hg
- (c) 0.96 atm
- (d) 1.02 atm
- 15. In an experiment during the analysis of a carbon compound, 145 l of  $H_2$  was collected at 760 mm Hg pressure and  $27^{o}C$  temperature. The mass of  $H_2$  is nearly

[MNR 1987]

- (a) 10 q
- (b) 12 q
- (c) 24 g
- (d) 6 g

- **16.** The volume of 1 g each of methane  $(CH_4)$ , ethane  $(C_2H_6)$ , propane  $(C_3H_8)$  and butane  $(C_4H_{10})$  was measured at 350 K and 1 atm. What is the volume of butane
  - (a)  $495 cm^3$
- (b)  $600 cm^3$
- (c)  $900 \text{ } cm^3$
- (d) 1700 cm<sup>3</sup>
- The ratio of the rate of diffusion of helium and methane under identical condition of pressure and temperature will be [IIT 2005]
  - (a) 4

(b) 2

(c) 1

(d) 0.5

At what temperature in the celsius scale, V (volume) of a certain mass of gas at  $27^{\circ}C$  will be doubled keeping the pressure constant[Orissa 1993]

- (a) 54°C
- (b) 327°C
- (c) 427°C
- (d) 527°C
- 19. Pressure of a mixture of 4 g of  $O_2$  and 2 g of  $H_2$  confined in a bulb of 1 litre at  $0^o C$  is [AIIMS 2000]
  - (a) 25.215 atm
- (b) 31.205 atm
- (c) 45.215 atm
- (d) 15.210 atm
- 20. If pressure becomes double at the same absolute temperature on 2  $L\,CO_2$ , then the volume of  $CO_2$  becomes

[AIIMS 1992]

- (a) 2 L
- (b) 4 L
- (c) 25 L
- (d) 1 L
- **21.** Volume of the air that will be expelled from a vessel of 300  $cm^3$  when it is heated from  $27^{\circ}C$  to  $37^{\circ}C$  at the same pressure will be
  - (a) 310  $cm^3$
- (b) 290  $cm^3$
- (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



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  $H_2$  and  $O_2$  each occupy

22.4 L of volume at  $0^{\circ}C$  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

constant 'a' greater is the

liquefaction of gas.

Reason : 'a' indirectly measures the

magnitude of attractive forces

between the molecules.

**6.** Assertion: Carbondioxide has greater value of

root mean square velocity  $\mu_{rms}$  than

carbon monoxide.

Reason :  $\mu_{rms}$  is directly proportional to

molar mass.

Reason

7. Assertion: 4.58 mm and  $0.0098 \,^{\circ} C$  is known to

be triple point of water.

At this pressure and temperature all the three states i.e., water, ice and

vapour exist simultaneously.

**8.** Assertion:  $1/4^{th}$  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 20

**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

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.

[AIIMS 2001]

**16.** Assertion: Effusion rate of oxygen is smaller

than nitrogen.

Reason: Molecular size of nitrogen is

smaller than oxygen. [AIIMS 2004]



#### Characteristics and Measurable properties of gases

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

# Ideal gas equation and Related gas laws

	1	С	2	d	3	a	4	а	5	а
	6	bc	7	a	8	а	9	С	10	d
	11	a	12	a	13	а	14	a	15	С
0	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	С	29	С	30	С
	31	а	32	а	33	b	34	b	35	b

250	Gaseous	state

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

21

b

22

# Kinetic molecular theory of gases and Molecular collisions

1	d	2	С	3	d	4	d	5	b
6	а	7	а	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	а	22	d	23	С	24	b	25	b
26	а	27	d	28	а	29	d	30	С
31	С								

# Molecular speeds

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

# Real gases and Vander waal's equation

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

# Critical state and Liquefaction of gases

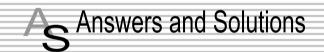
1	а	2	b	3	d	4	С	5	d
6	d	7	С	8	С	9	abc	10	С
11	С								

# **Critical Thinking Questions**

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

#### **Assertion & Reason**

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



# Characteristics and Measurable properties of gases

- (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  $0^{\circ} C \& 4.7 \ mm$  pressure.
- **3.** (b) It is characteristic of gases *i.e.* Thermal energy >> molecular attraction
- 4. (a) In gases, molecular attraction is very less and intermolecular spaces are large hence kinetic energy of gases is highest.
- 5. (c) Gases and liquids, both can flow and posses viscosity.
- 7. (a) Newton is unit of force.
- **8.** (b)  $\frac{C^o}{5} = \frac{F^o 32}{9}$
- 9. (c)  $1L = 10^{-3}m^3 = 10^3 cm^3 = 1 dm^3 = 10^3 ml$ .
- **10.** (a) 1  $atm = 10^6$  dynes cm<sup>-2</sup>
- 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)  $0^{\circ} C$  is equivalent to  $273^{\circ} Ki.e.$  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

$$P=100\ kPa$$
 ,  $P_1=?$  ,  $V=1\ litre$  ,  $V_1=4litre$ 

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

- 1. (c) Boyle's law is  $V \propto \frac{1}{P}$  at constant T
- 2. (d) According to Boyle's law  $V \propto \frac{1}{P}$

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

- 3. (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.
- 4. (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.

7. (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

11. (a) 
$$T_1 = 273^{\circ}C = 273 + 273^{\circ}K = 546^{\circ}K$$
  
 $T_2 = 0^{\circ}C = 273 + 0^{\circ}C = 273^{\circ}K$   
 $P_1 = 1$ ;  $P_2 = ?$   
According to Gay-Lussac's law  
 $P_1 = P_2$  .  $P_1 = P_1 = 1 \times 273^{\circ}K$  atm: 1

 $: (V_2 - V_1) = \Delta V = V_0 \alpha (t_2 - t_1)$ 

$$\frac{P_1}{T_1} = \frac{P_2}{T_2} \therefore P_2 = \frac{P_1 T_2}{T_1} = \frac{1 \times 273^{\circ} K}{546^{\circ} K} \text{ atm; } \frac{1}{2} \text{ atm.}$$
**12.** (a)  $V_t = V_o (1 + \alpha_v t)$ 

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.

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

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

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

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

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

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

 $\frac{m_1}{m_2} = \frac{V_2}{V_1} \therefore m_1 < m_2$  is curve for  $V_2$  has a greater

slope than for V<sub>1</sub>

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

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

**18.** (c) At constant *V* of a definite mass

At constant V of a definite mass 
$$\frac{P_1}{T_1} = \frac{P_2}{T_2} \therefore \frac{P_1}{P_2} = \frac{300}{600} = \frac{1}{2} \text{ i.e. pressure increases}$$

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.

**19.** (a) Avogadro number =  $6.0224 \times 10^{23}$ 

**20.** (b) Compressibility = 
$$\frac{PV}{nRT}$$
 < 1 at STP (as given)

$$nRT > PV$$
  
 $N \times 0.0821 \times 273 > 1V_m$   
 $22.41 \ litres > V_m$   
 $R = 0.821$   
 $T = 273^{\circ}K$   
 $P = 1$   
 $n = 1$ 

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

**22.** (c) 
$$PV = nRT$$

$$R = \frac{PV}{nT} = \text{ litre . atm. } K^{-1} \text{ mole}^{-1}$$

**23.** (d)  $(atm. K^{-1} mol^{-1})$  is not a unit of R

24. (a) 
$$8.31 J.K^{-1} mol^{-1}$$
  
 $1 cal = 4.2 J.$   
 $\therefore \frac{8.31}{4.2} cal.K^{-1} mol^{-1} = 1.987 cal K^{-1} mol^{-1}$ 

**30.** (c) 
$$PV = nRT : \frac{n}{V} = \frac{P}{RT}$$

31. (a) 
$$P = \frac{nRT}{V} = \frac{2 \times 0.0821 \times 546}{44.81} = 2 \text{ atm.}$$

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

at STP 
$$n_1$$
 = one mole, at  $T = 273^{\circ} + 30^{\circ} = 303^{\circ} K$   
 $P_1 = 1$  atm.  $P_2 = 1$  atm  $P_3 = 1$  atm  $P_4 = 1$  atm  $P_2 = 1$  atm  $P_3 = 1$  atm  $P_4 =$ 

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

**34.** (b) 
$$V_2 = \frac{P_1 V_1 T_2}{P_2 T_1} \Rightarrow P_1 = P$$
 ;  $T_1 = 273^{\circ} K$ 

$$P_2 = \frac{3}{2} P \text{ ; } T_2 = T_1 + \frac{T_1}{3} = \frac{4}{3} \times 273^{\circ} K$$

$$V_2 = \frac{2P}{3P} \times \frac{4}{3} \times \frac{273}{273} \times 100 \text{ } cc = \frac{800}{9} \text{ } cc = 88.888 \text{ } cc$$

$$= 88.9 \text{ } cc$$

**35.** (b) 
$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T} : \frac{P_1V_1}{P_2V_2} = \frac{T_1}{T_2}$$

**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) 
$$n = \frac{PV}{RT} = \frac{m}{M}$$
  
 $m = \frac{MPV}{RT} = \frac{34 \times 2 \times 100}{0.082 \times 293} = 282.4 \text{ gm}$ 

**39.** (c) 
$$V_2 = \frac{P_1 V_1}{T_1} \frac{T_1}{P_2} = \frac{760}{600} \times \frac{546}{273} \times 273 = 691.6 ml.$$

**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) 
$$V_2 = \frac{P_1}{P_2} \frac{T_2}{T_1} . V_1 = \frac{1}{0.5} \times \frac{250}{300} \times 12000 \ lit. = 20000 \ lit.$$

**42.** (c) At constant pressure

$$V \propto nT \propto \frac{m}{M} T$$

$$\frac{V_1}{V_2} = \frac{m_1 T_1}{m_2 T_2} \therefore \frac{T_1}{T_2} = \frac{V_1}{m_1} \times \frac{m_2}{V_2} = \frac{d_2}{d_1} \Rightarrow \frac{300^{\circ} K}{T_2} = \frac{0.75 d}{d}$$

$$T_2 = \frac{300}{0.75} = 400^{\circ} K$$

**43.** (c) 
$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$
 :  $T_2 = \frac{P_2 V_2}{P_1 V_1} \times T_1 = \frac{740}{740} \times \frac{80}{100} \times 300$   
=  $240^{\circ} K = -33^{\circ} C$ 

- **47.** (b) Because  $H_2$  &  $Cl_2$  gases may react with each other to produce HCl gas hence Dalton's law is not applicable.
- **48.** (d) Because  $HCl \& NH_3$  gases may react to produce  $NH_4Cl$  gas. Dalton's Law is applicable for non reacting gas mixtures.
- **49.** (b)  $NH_3$  and HCl & HBr is a reacting gas mixture to 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  $=\frac{\frac{m}{4}}{11m}=\frac{10}{11}$

Partial pressure due to lighter gas =  $P_o \times \frac{10}{11}$ =  $1.1 \times \frac{10}{11} = 1$  atm.

**52.** (a) m. wt. of  $NH_3 = 17$ ; m.wt. of  $N_2 = 28$  m.wt. of  $CO_2 = 44$ ; m.wt. of  $O_2 = 32$ beacuse  $NH_3$  is lightest gas out of these gases  $r \propto \frac{1}{\sqrt{\text{Molecular Weight}}}$ 

55. (a) 
$$\frac{r_g}{r_{He}} = \sqrt{\frac{M_{He}}{M_g}} : M_g = M_{He} \cdot \frac{r^2_{He}}{r^2_g} = \frac{4}{(1.4)^2} = \frac{4}{1.96} = 2$$

[Note: 1.4 =  $\sqrt{2}$ ]

**56.** (a) 
$$r_g = \frac{1}{5} . r_{H_2}$$

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

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

**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) 
$$r_O = r_H \sqrt{\frac{d_H}{d_O}} = 1\sqrt{\frac{0.09}{1.44}} = \sqrt{\frac{1}{16}} = \frac{1}{4}$$

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

- 63. (c) Gases may be separated by this process because of different rates of diffusion due to difference in their densities.
- **64.** (b)  $NH_4Cl$  ring will first formed near the HCl bottle because rate of diffusion of  $NH_3$  is more than that of HCl because  $M_{NH_3}: M_{HCl} = 17:36.5)$ . SO  $NH_3$  will reach first to the HCl bottle & will react there with HCl to form  $NH_4Cl$  ring
- **65.** (d) Because both *NO* and  $C_2H_6$  have same molecular weights  $\left\lfloor M_{NO} = M_{C_2H_6} = 30 \right\rfloor$  and rate of diffusion  $\infty$  molecular weight.

**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) 
$$r_H = \frac{2gm}{10 \text{ min}}$$
 if  $r_O = \frac{xgm}{10 \text{ min}}$ 

$$r_O = r_H \sqrt{\frac{M_{H_2}}{M_{O_2}}} = \frac{2}{10} \sqrt{\frac{2}{32}}$$

$$\frac{x}{10} = \frac{2}{10 \times 4} = \frac{1}{2} gm. = .5 gm$$

**69.** (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) 
$$r \propto \frac{1}{\sqrt{M}}$$
  $\therefore r = \frac{Volume \ effused}{time \ taken} = \frac{V}{t}$ 

$$\frac{V}{t} \propto \frac{1}{\sqrt{M}} \therefore \text{for same volumes ($V$ constant)}$$

$$t \propto \sqrt{M} \therefore \frac{t_1}{t_2} = \sqrt{\frac{M_1}{M_2}}$$

$$t_{He} = t_{H_2} \sqrt{\frac{M_{He}}{M_H}} = 5\sqrt{\frac{4}{2}} = 5\sqrt{2}s.$$

$$t_{O_2} = t = 5\sqrt{\frac{32}{2}} = 20 \, s$$

$$t_{CO} = 5\sqrt{\frac{28}{2}} = 5\sqrt{14}s$$
;  $t_{CO_2} = 5\sqrt{\frac{44}{2}} = 5\sqrt{22}s$ 

71. (c) 
$$\frac{r_{N_2}}{r_{SO_2}} = \frac{V_{rms}N_2}{V_{rms}SO_2} = \sqrt{\frac{T_{N_2}}{T_{SO_2}} \cdot \frac{M_{SO_2}}{M_{N_2}}} = \sqrt{\frac{T_{N_2}}{323} \times \frac{64}{28}}$$

$$1.625 = \sqrt{\frac{T_{N_2}}{323} \cdot \frac{16}{7}}$$

$$T_{N_2} = \frac{(1.625)^2 \times 323 \times 7}{16} = 373^{\circ} K$$

**72.** (a) 
$$C + H_2O \rightarrow CO_{(g)} + H_{2(g)}$$

 $12\,gm \rightarrow 1mol + 1mol$ 

12 gm C produces 2mole of gases (1mole CO & 1 mole of  $H_2$ )

∴48 gm C may produce 
$$\frac{48}{12} \times 2 = 4 \times 2 = 8$$
 mole

 $=22.4\times8 L \text{ gases} = 179.2 L \text{ gas.}$ 

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.** (c) 
$$PV = nRT$$

$$P = \frac{n}{V}RT \quad \because \frac{n}{V} = C \implies P = CRT$$

$$T = \frac{P}{CR} = \frac{1}{1 \times .821} = 12^{\circ} K$$

**75.** (a) 
$$6.02 \times 10^{22}$$
 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  $O_2$ 

$$=(28\times0.1)+(2\times0.1)+(32\times0.1) = 6.2\,gm$$

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  $SO_2$  and  $Cl_2$  are reacted chemically and forms  $SO_2Cl_2$ . That is why mixture of these gases is not applicable for Dalton's law.

**86.** (d) According to Boyle's law 
$$P_1V_1 = P_2V_2 \implies P_1 \times 60 = 720 \times 100$$
  $P_1 = \frac{720 \times 100}{60} = 1200 \, mm$ 

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

that is why  $H_2$  gas diffuse first

88. (a) Solution level will rise, due to absorption of  ${\it CO}_2$  by sodium hydroxide.

$$2NaOH + CO_2 \rightarrow Na_2CO_3 + H_2O$$

**89.** (c) 
$$CaCo_3 \xrightarrow{} \hat{} CaO + CO_2 \uparrow \\ (40+12+16\times3)=100 \text{ gm} \xrightarrow{} CaO + CO_2 \uparrow \\ 22.4 \text{ litre}$$

: At S.T.P.  $100 g CaCO_3$  produce= 22.4 litre of  $CO_2$ 

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

**90.** (c) The density of gas 
$$=\frac{Molecular\ wt.\ Of\ Metal}{Volume}$$

$$= 2gmlitre^{-1}$$

**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  $= 6.023 \times 10^{23}$ 

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  $= \frac{6.023 \times 10^{23}}{22.4} \times 15 = 4.03 \times 10^{23} \text{ molecules.}$ 

**93.** (a) 22.4
$$l$$
  $O_2$  at S.T.P. =  $32 gm$  of  $O_2$   
 $1l$   $O_2$  at S.T.P. =  $\frac{32}{224} = 1.43 gm$  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

$$\frac{r_1}{r_2} = \sqrt{\frac{M_2}{M_1}} = \sqrt{\frac{4}{2}} = \sqrt{2} = 1.4$$
;  $r_1 = 1.4m$ 

**96.** (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:

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

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,  $P_2 = 857.14 mm$ 

98. (b) According to gas law

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

$$\frac{n_A}{n_B} = \frac{\frac{P_1 V_1}{R T_1}}{\frac{P_2 V_2}{R T_2}}; \quad \frac{n_A}{n_B} = \frac{P_1 V_1}{T_1} \times \frac{T_2}{P_2 V_2}$$

$$\frac{n_A}{n_B} = \frac{2P \times 2V}{2T} \times \frac{T}{PV} \; ; \quad \frac{n_A}{n_B} = \frac{2}{1}$$

**99.** (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  $NO_2 = \frac{20.10}{46} = 0.437$ 

100. (a) Given that

 $P_1 = 76cm$  of Hg (Initial pressure at N.T.P.)

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

According to Boyle's law

$$P_1V_1 = P_2V_2$$
;  $76 \times 5 = P_2 \times 35$ 

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

**101.** (c) Given initial volume  $(V_1) = 300\,cc$ , initial temperature  $(T_1) = 27^{\circ}\,C = 300\,K$ , initial pressure  $(P_1) = 620\,mm$ , final temperature  $T_2 = 47^{\circ}\,C = 320\,K$  and final pressure  $(P_2) = 640\,mm$ . We know from the general gas equation

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2} = \frac{620 \times 300}{300} = \frac{640 \times V_2}{320} \implies V_2 = 310 cc$$

**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  $V_2 = 4 dm^3$ 

**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)  $P_1 = P, V_1 = V, T_1 = 273 + 75 = 348 K$ 

$$P_2 = 2P, V_2 = \frac{85}{100}, T_2 = ?$$

$$\frac{P \times V}{398} = \frac{2P \times 85 \, V}{T_2 \times 100} \Rightarrow T_2 = \frac{348 \times 2 \times 85}{100}$$

$$T_2 = 591.6K = 318.6^{\circ} C$$

**106.** (a) Boyle's law -PV = constant On differentiating the equation,

$$d(PV) = d(C) \implies PdV + VdP = 0$$

$$\Rightarrow VdP = -PdV \Rightarrow \frac{dP}{P} = -\frac{dV}{V}$$
.

# 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  $(V = 1) \Rightarrow P = \frac{2}{3} E$ 

8. (c) Tr.  $K.E. = \frac{3RT}{2}$  it means that the Translational Kinetic energy of Ideal gas depends upon

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.

temperature only.

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

K.E. = 37

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.

**15.** (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)

- **16.** (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)  $PV = nRT = \frac{m}{M}RT$  $\therefore \frac{m}{V} = \frac{PM}{RT} = \text{density}$
- **19.** (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
- 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.

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

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

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

- 22. (d) Density =  $\frac{M}{V}$   $d = \frac{v.d \times 2}{V} \qquad (M = V.d \times 2)$   $V.d = \frac{d \times V}{2}$   $V.d = \frac{0.00130 \times 22400}{2} = 14.56 \text{ gm}^{-1}$
- 23. (c) Volume of steam =  $1lt = 10^3 cm^3$   $\therefore m = d.V$   $\therefore$  mass of  $10^3 cm^3$  steam = density × Volume =  $\frac{0.0006 gm}{cm^3} \times 10^3 cm^3 = 0.6 gm$

Actual volume occupied by  $H_2O$  molecules is equal to volume of water of same mass

- $\therefore$  Actual volume of  $H_2O$  molecules in 6gm steam
- = mass of steam/density of water =  $0.6 \text{ gm} / 1 \text{ qm/cm}^3 \Rightarrow 0.6 \text{ cm}^3$
- **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)  $d \propto M \Rightarrow \frac{d_1}{d_2} = \frac{M_1}{M_2}; \frac{3d}{d} = \frac{M}{M_2}; M_2 = \frac{M}{3}.$

# Molecular speeds

- 1. (a)  $V_{ms} = \sqrt{\frac{3RT}{M}}, V_{av} = \sqrt{\frac{8RT}{\pi M}}; \frac{V_{ms}}{V_{av}} = \sqrt{\frac{3\pi}{8}}$  $= \sqrt{\frac{66}{56}} \Rightarrow \frac{1.086}{1}$
- 3. (d) most probable velocity: mean velocity:  $V_{rms} = \sqrt{\frac{2RT}{M}} : \sqrt{\frac{8RT}{\pi M}} : \sqrt{\frac{3RT}{M}} = \sqrt{2} : \sqrt{\frac{8}{\pi}} : \sqrt{3}$
- 4. (d)  $V_{ms} = \sqrt{\frac{3RT}{M}} : V_{ms} \propto \frac{1}{\sqrt{M}}$  at same T

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

- 5. (d)  $\frac{U_{SO_2}}{U_{He}} = \frac{1}{2} = \sqrt{\frac{M_{He}}{M_{SO_2}} \frac{T_{SO_2}}{T_{He}}} = \sqrt{\frac{4}{64} \cdot \frac{T_{SO_2}}{300}}$ =  $\frac{4}{64} \cdot \frac{T_{SO_2}}{300} = \frac{1}{4}$ ;  $T_{SO_2} = 1200^{\circ} K$
- **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  $= \frac{3}{2} KT = \frac{3}{2} \times 1.38 \times 10^{-23} \times 300 J = 6.17 \times 10^{-21} J$
- **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) 
$$V_{rms} = \sqrt{\frac{3KT}{Molecular weight}}$$
 i.e.  $V_{rms} \propto \frac{1}{\sqrt{m}} \propto (m)^{-\frac{1}{2}}$ 

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

$$\frac{U_1^2}{U_2^2} = \frac{m_2}{m_1} \qquad \therefore m_1 U_1^2 = m_2 U_2^2$$

**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) 
$$U = \sqrt{\frac{3RT}{M}} = \sqrt{\frac{3PV}{M}} = \sqrt{\frac{3P}{d}} : U \propto \frac{1}{\sqrt{d}}$$

**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  $V_1 = V$ ,  $T_1 = 300 \, K$ ,  $V_2 = 2V$ ,  $T_2 = ?$   
 $\frac{V_1}{V_2} = \sqrt{\frac{T_1}{T_2}}$ ;  $\left(\frac{V}{2V}\right)^2 = \frac{300}{T_2} \Rightarrow T_2 = 300 \times 4 = 1200 \, K$ 

**24.** (b) 
$$\left(P - \frac{n^2 a}{V^2}\right) (V - nb) = nRT$$
  
 $\left(P - \frac{(2)^2 \times 4.17}{(5)^2}\right) (5 - 2 \times .03711) = 2 \times .0821 \times 300$   
 $P = \frac{2 \times .0821 \times 300}{5 - 2 \times .03711} - \frac{4.7 \times 2^2}{5^2} \Rightarrow 10 - 0.66 = 9.33 atm$ 

**26.** (a) Average speed : most probable speed 
$$\sqrt{\frac{8RT}{\pi M}}: \sqrt{\frac{2RT}{M}} \implies \sqrt{\frac{8}{\pi}}: \sqrt{2} \implies 1.128: 1.$$

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

$$\sqrt{30^2 R} = \sqrt{\frac{3RT}{M}} \implies 30 \times 30 R = \frac{3R \times 300}{M}$$

$$\Rightarrow M = \frac{3 \times 300}{30 \times 30} 1 \text{ gm} = 0.001 \text{ kg}.$$

# 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.

3. (c) 
$$Z = \frac{PV}{RT}$$
 : for ideal gas  $PV = RT$  so  $Z = 1$ 

**12.** (b) 
$$Z = \frac{PV}{RT}$$
; for ideal gas  $PV = RT$ ; so  $Z = 1$ 

- **13.** (b) Ideal gas has no attractive force between the particles
- 14. (d) PV = nRT is a ideal gas equation it is allowed when the temperature is high and pressure is low.
- **16.** (b) At Boyle temperature real gas is changed into ideal gas
- 17. (a) When pressure is low  $\left[P + \frac{a}{V^2}\right](V b) = RT$ 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)
  20. (a) Vander waal's constant for volume correction b is the measure 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  $(P + 36.048)(0.2072) = 24.63$   
 $\Rightarrow P + 36.048 = 118.87 \Rightarrow P = 82.82 \text{ atm.}$ 

# Critical state and Liquefaction of gases

- (b) A diabatic demagnetisation is a technique of liquefaction of gases in which temperature is reduced.
- 3. (d) An ideal gas can't be liquefied because molecules of ideal gas have not force of attraction between them.
- **4.** (c) At above critical temperature, substances are existing in gaseous state, since gas cannot be liquefied above it.

- (d) Absence of inter molecular attraction ideal gas 5. cannot be liquefied at any volume of P and T.
- 6. (d) For Z gas of given gases, critical temperature is highest

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

- (c) Value of constant a is greater than other for 7.  $NH_3$  that's why  $NH_3$  can be most easily liquefied.
- (c) The temperature below which the gas can be 8. liquefied by the application of pressure alone is called critical temperature.
- (c)  $d = \frac{M}{V} \implies M = d \times V$ 10.  $M = 2.68 \times 22.4$  at N.T.P. (: V = 22.41)  $M = 60.03 \ gm$

m. wt of COS = 12 + 16 + 32 = 60

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

1ml 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  $O_2$ 

# **Critical Thinking Questions**

(a) Average kinetic energy ∞ (T Kelvin) 1.

(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  
= 
$$70 \times 2 = 140 \Rightarrow x = \frac{m.wt.}{wt.of[CO]} = \frac{140}{[12+16]} = 5$$

- (c) Gas deviate from ideal gas behaviour to real 3. gas (according to Vander Waal's at low temperature and high pressure)
- (c) At same temperature and pressure, equal 4. volumes have equal number of molecules. If 1lit. of oxygen consists N molecules then at same temperature and pressure 1 lit of SO2 will consists N molecules. So 2 lit. of  $SO_2$  will contain 2N molecules.
- $V_{av}:V_{rms}:V_{most\ probable}=$ (a) 5.

$$\sqrt{\frac{8RT}{\pi M}} : \sqrt{\frac{3RT}{M}} : \sqrt{\frac{2RT}{M}}$$

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

- (d)  $\frac{C_p}{C}$  ratio for diatomic gases is 1.40 at lower 6. temperature & 1.29 at higher temperature so
- the answer is 'd'. (b) PV = nRT (For ideal gas) 7.

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

- (b) If Z < 1 then molar volume is less than 22.4 L
- (a)  $r_{O_2} = \frac{x}{18} mole / sec \implies r_g = \frac{x}{45} mol / sec$

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

- **10.** (a)  $r_{SO_2}: r_{O_2}: r_{CH_4} = \frac{1}{\sqrt{M_{SO_2}}}: \frac{1}{\sqrt{O_2}}: \frac{1}{\sqrt{CH_4}}$  $=\frac{1}{\sqrt{64\cdot 32\cdot 16}}=\frac{1}{\sqrt{4\cdot 2\cdot 1}}$  $\frac{1}{2}:\frac{1}{\sqrt{2}}:\frac{1}{1}; \frac{2}{2}:\frac{2}{\sqrt{2}}:\frac{2}{1}; 1:\sqrt{2}:2$
- square speed =  $\left[\frac{n_1c_1^2 + n_2c_2^2 + n_3c_3^2 + ...}{n_1 + n_2 + n_3 + ...}\right]^{1/2}$ .
- (b) If 40 ml  $O_2$  will diffuse in t min. then.  $r_H = \frac{50}{20} \implies r_O = r_H 2 \sqrt{\frac{M_{H_2}}{M_{O_2}}} = \frac{50}{20} \sqrt{\frac{2}{32}} = \frac{50}{20} \cdot \frac{1}{4}$  $\frac{40}{t} = \frac{50}{80} : t = \frac{40 \times 80}{50} = 60 \quad min.$
- **13.** (b)  $\frac{V_{av}CH_4}{V_{ab}O_2} = \sqrt{\frac{T_{CH_4}}{T_{O_2}} \cdot \frac{M_{O_2}}{M_{CH_4}}} = 1$  $\frac{T_{CH_4}}{300} \cdot \frac{32}{16} = 1$ ;  $T_{CH_4} = 150^{\circ} K$
- **14.** (c) Pressure of  $O_2$  (dry) = 751-21 = 730 mm Hg  $=\frac{730}{760}=0.96 atm$

**15.** (b) 
$$PV = nRT$$
,  $n = \frac{PV}{RT} = \frac{1 \times 145}{0.082 \times 3} = 5.8 \approx 6 \, mole$ .

**16.** (a) 
$$V = \frac{nRT}{P} = \frac{m}{M} \cdot \frac{RT}{P} = \frac{1}{58} \times \frac{0.82 \times 350}{1} = 0.495 \, lit.$$
  
= 495 cm<sup>3</sup>

17. (b) 
$$\frac{r_{He}}{r_{CH_4}} = \sqrt{\frac{M_{CH_4}}{M_{He}}} = \sqrt{\frac{16}{4}} = 2$$

**18.** (b) 
$$\frac{V_1}{V_2} = \frac{T_1}{T_2} :: T_2 = \frac{T_1 V_2}{V_1} = 300^{\circ} K, \frac{2V}{V} = 600^{\circ} K$$

$$T_2 = 600^{\circ} K = (600 - 273)^{\circ} C = 327^{\circ} C$$

**19.** (a) no. of moles of  $O_2 = \frac{4}{32} = 0.125$ no. of moles of  $H_2 = \frac{2}{3} = 1$ total no. of moles = 1+0.125 = 1.125 $P = \frac{nRT}{V} = \frac{1.125 \times 0.082 \times 273}{1} = 25.184 \text{ atm.}$ 

**20.** (d) 
$$\frac{P_1}{P_2} = \frac{1}{2}$$
,  $\therefore \frac{V_1}{V_2} = \frac{P_2}{P_2} = \frac{2}{1}$ 

$$\frac{2L}{V_2} = \frac{2}{1}$$
;  $V_2 = 1L$ 

**21.** (c) 
$$\frac{V_2}{V_1} = \frac{T_2}{T_1}$$

$$V_2 = \frac{T_2}{T_1} V_1 = \frac{310^{\circ} K}{300^{\circ} K} \times 300 \text{ cm}^3 = 310 \text{ cm}^3$$

**22.** (c) 
$$V_2 = \frac{T_2}{T_1} \cdot V_1 = \frac{270^{\circ} K}{300^{\circ} K} \times 300 ml = 270 ml$$

#### **Assertion & Reason**

- 1. (c) Pressure is inversly proportional to volume (Boyle's law).  $p\alpha \frac{1}{V}$  (n, T constant).
- 2. (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.
- 3. (a) At a given temperature and pressure the volume of a gas is directly proportional to the amount of gas Von (P and T constant).
- 4. (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)  $P_{Total} = P_1 + P_2 + P_3 \dots$  (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)  $\mu_{rms} = \sqrt{\frac{3RT}{M}}$  is inversly related to molecular mass. Therefore,  $\mu_{rms}(CO) > \mu_{rms}(CO_2)$ .
- **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  $\frac{V_1}{V_2} = \frac{300}{400} = \frac{3}{4}$  : air expelled  $\Rightarrow 1 \frac{3}{4} = \frac{1}{4}$
- **9.** (a) In case of  $H_2$ , 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 molecules increases and by this increase the pressure of gas is also increases.
- 12. (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.
- **14.** (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.