



Compressibility, Boyle's Temperature

Course on States of Matter for Class XI

9

$$\frac{6 \times 10^{20}}{6 \times 10^{23}} = 10^{-3}$$

S-II

$$KE_{\text{mol}} = \frac{3}{2} \times \overset{4}{\cancel{8}} \times 3 \omega$$

$$= 36 \omega$$

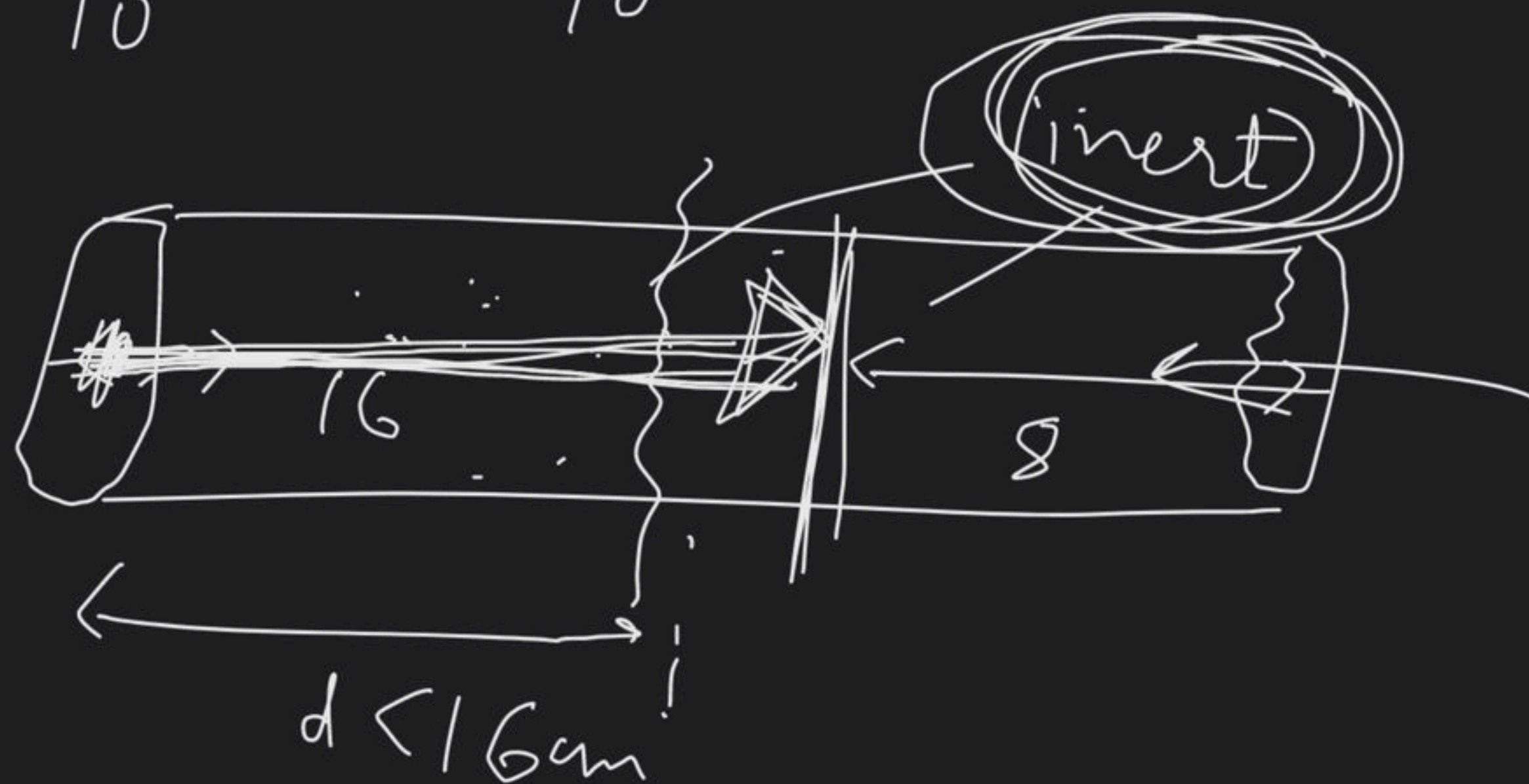
$$= \underline{3.6}$$

$$P^2/T^{3/2}$$

$$\lambda = \frac{\hbar T}{\sqrt{2} \pi \sigma^2 P}$$

$$2.6 \times 10^{-5} = \frac{8.314}{N_A} \times \frac{300}{\sqrt{2} \pi \sigma^2 P}$$

X Y
 10 40
 $\sigma_y = \sigma_y$



1 Sec

Z_{11}

$$\underbrace{Z_{12}}_{\text{circled}} = \pi \sigma^2 \sqrt{\frac{8RT}{\pi} \left(\underbrace{M_1}_{\text{circled}} + \frac{1}{M_2} \right)} \times N_{\text{inert}}^x$$

diffusion coefficient $\propto \lambda V_{\text{avg}}$

$$\propto \frac{T}{P} \times \sqrt{T}$$

$$\propto \frac{T^{3/2}}{P}$$

$$\frac{(4)^{3/2}}{2} = \frac{8}{2} = 4$$

$$P = P_{\text{obs}} \oplus \underline{P_{\text{corr}}}$$

Consider a molecule 'B' which is about to collide the wall of the container.

This molecule will experience a retarding force towards the centre of container due to which its speed decreases.

Hence it exerts less force than
it would have if there were no forces
of attraction.

$$\left(p + \frac{an^2}{V^2} \right) (V - nb) = nRT$$

↑
Pressure of
gas

↑
Vol. of gas
= V_{cont}

↑

a, b are temperature independent
but gas dependent
Vander Waal's const

find P of 1 mol real gas in a 2 lit container at 300 K. Given $a = 2 \text{ atm lit}^2 \text{ mol}^{-2}$

$R = 0.08 \text{ atm lit/mol/K}$ $b = 0.4 \text{ lit/mol}$

$$P \times 2 = 1 \times 0.08 \times 300$$

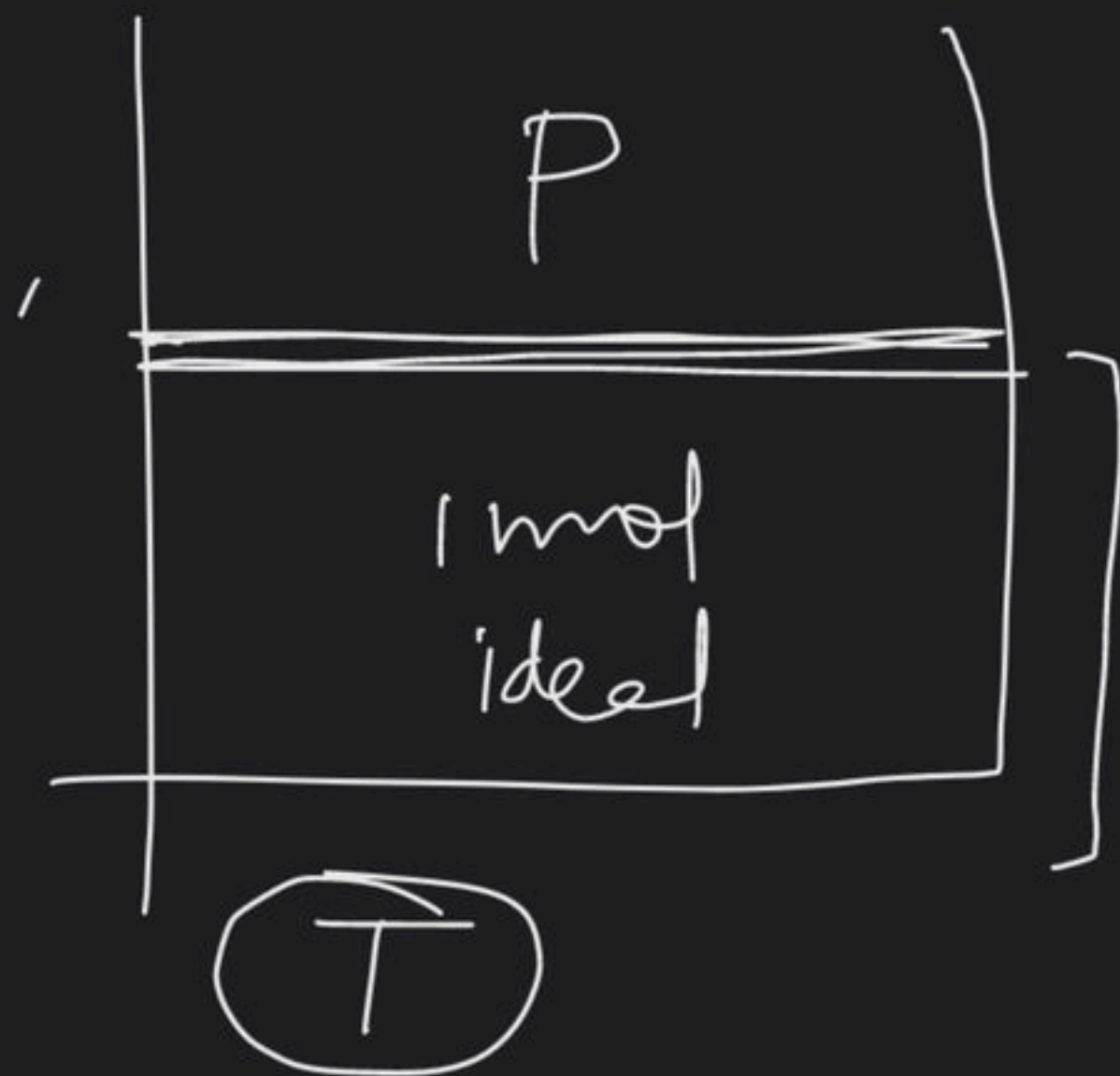
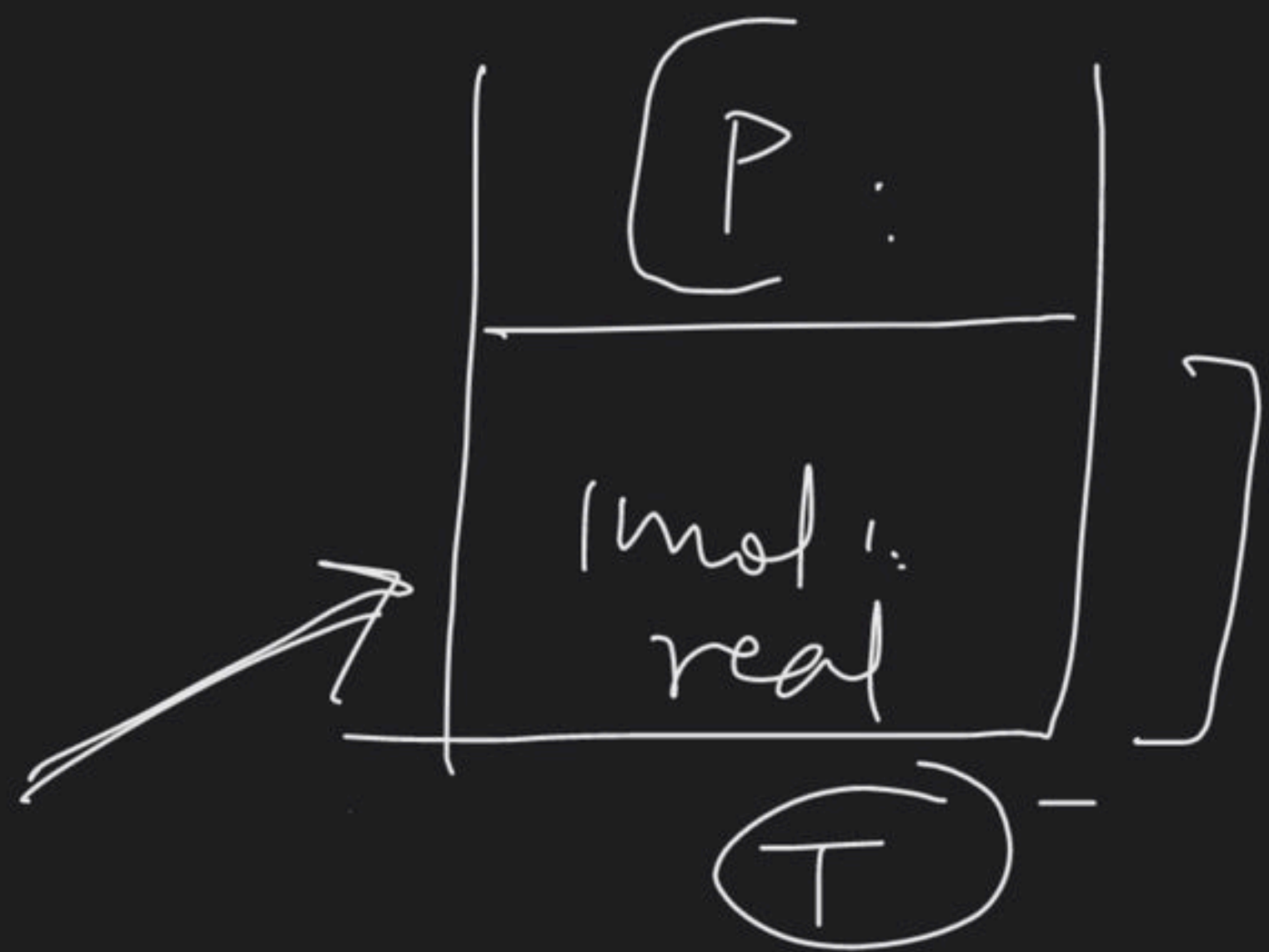
$$\underline{\underline{P = 12}}$$

- A) 12
- B) 14.5
- C) 15.5
- D) None

Compressibility factor (Z)

$$Z = \frac{V_{\text{real}}}{V_{\text{ideal}}} = \frac{\text{Vol of real gas}}{(\text{Vol. of ideal gas at same } T, P)} = \frac{PV}{nRT}$$

$\downarrow \downarrow$
 $P V$
 $\uparrow \uparrow$
 $n R T$



$$V = \frac{nRT}{P}$$

$$Z = \frac{PV}{nRT} = \frac{V_{\text{real}}}{V_{\text{ideal}}}$$

$$\underline{a = 0.5}$$

$$PV = Z n R T$$

$$\left(p + \frac{an^2}{V^2} \right) (V - nb) = nRT$$

Z vs P (At const n & T)

$$V_m = \frac{V}{n}$$

molar volume

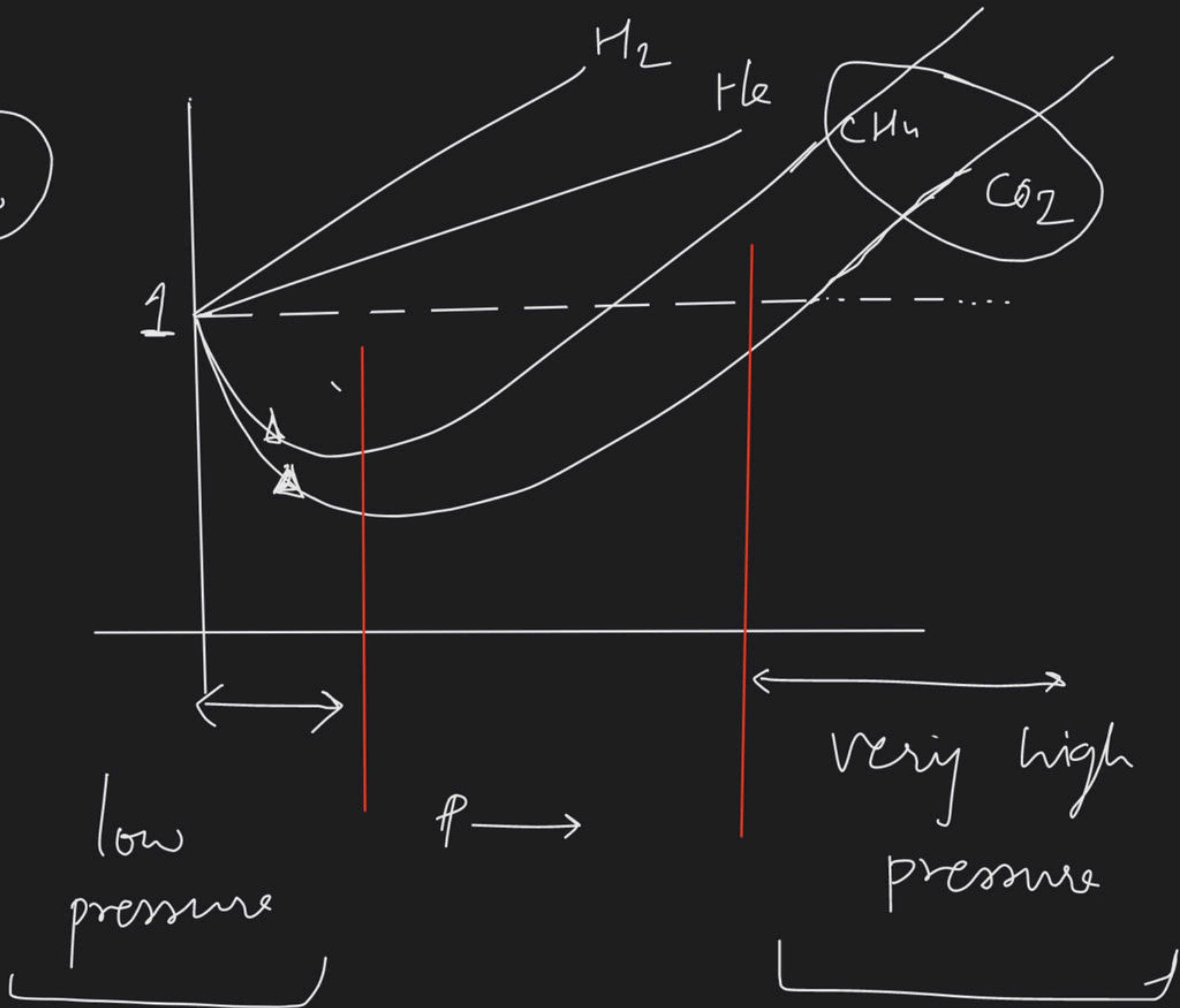
$$Z = \frac{PV}{nRT} = \frac{PV_m}{RT}$$

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

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

Z

At $T = 298\text{ K}$



Q.2 If Z is the compressibility factor, van der Waals' equation at low pressure can be written as : **[JEE-MAINS-2014]**

(1) $Z = 1 - \frac{Pb}{RT}$

(2) $Z = 1 + \frac{Pb}{RT}$

(3) $Z = 1 + \frac{Pb}{RT}$

(4) $Z = 1 - \frac{a}{V_m RT}$

Q.3 The compressibility factor for a real gas at high pressure is :- **[AIIEEE-2012]**

(1) $1 - \frac{Pb}{RT}$

(2) $1 + \frac{PT}{Pb}$

(3) 1

(4) $1 + \frac{Pb}{RT}$

Case - I

In low pressure region

$$\left(\underset{\substack{\uparrow \\ \text{low}}}{P} + \frac{a}{\underset{\substack{\uparrow \\ \text{low}}}{V_m^2}} \right) \left(\underset{\substack{\uparrow \\ \text{high}}}{V_m} - \underset{\substack{\uparrow \\ \text{low}}}{b} \right) = RT$$

$$Z = 1 - \frac{a}{V_m RT}$$

$$\textcircled{Z} = \frac{PV_m}{RT} = 1 - \frac{a}{\textcircled{V_m RT}}$$

Case-II

At very high pressure $\therefore \rightarrow$

A+ P
high

V
low

A+ very high

low

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

↑ ↑ ↑
very high high low low

$$\frac{PV_m}{RT} - \frac{Pb}{RT} = \frac{RT}{RT}$$

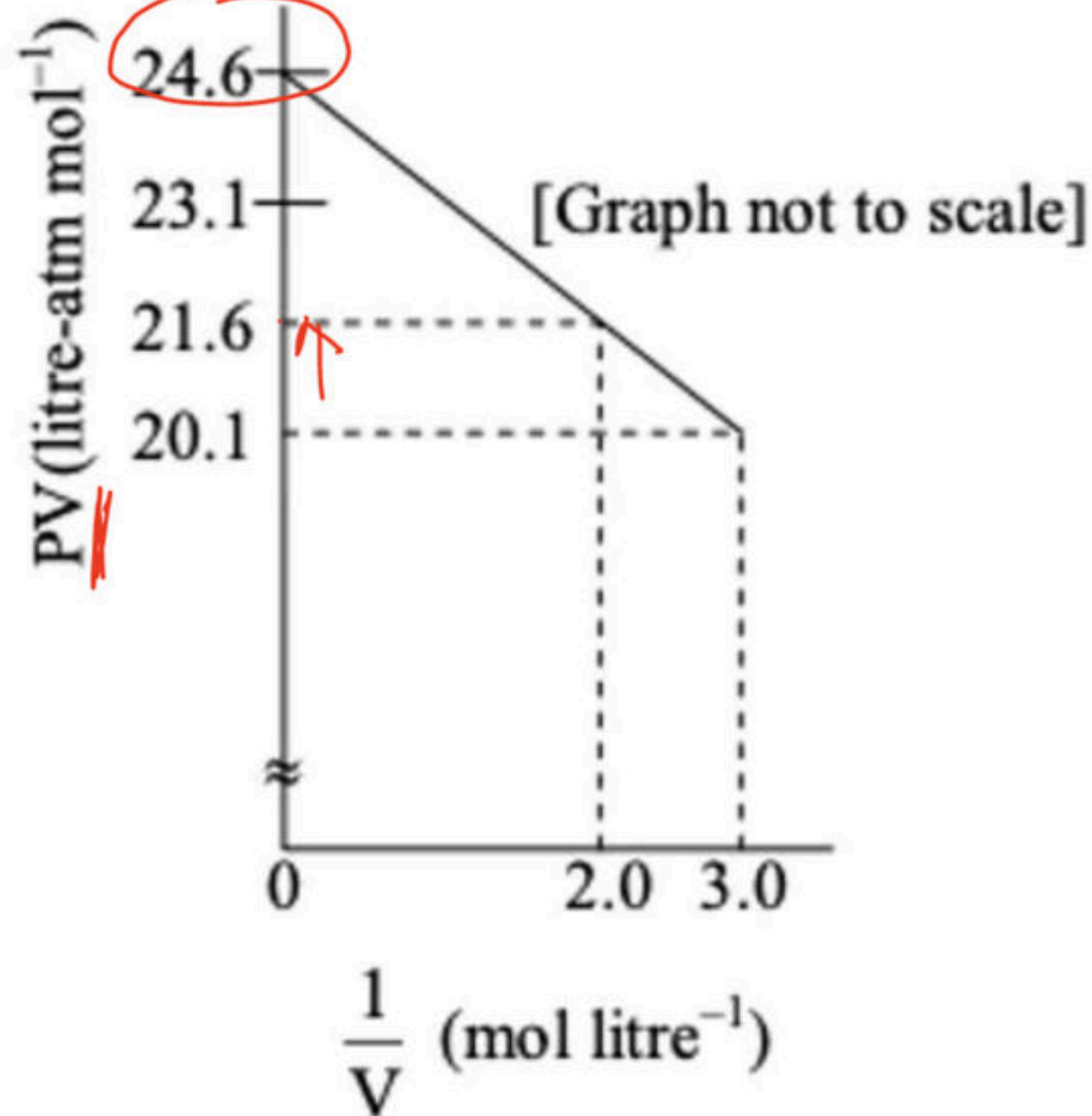
$$Z = 1 + \frac{Pb}{RT}$$

Q.13 For one mole of a van der Waals' gas when $b = 0$ and $T = 300\text{ K}$, the PV vs. $1/V$ plot is shown below. The value of the van der Waals' constant a ($\text{atm. litre}^2 \text{ mol}^{-2}$) is

[JEE 2012]

$$\frac{PV_m}{RT} = 1 - \frac{a}{V_m RT}$$

$$\textcircled{PV_m} = RT - \textcircled{\frac{a}{V_m}}$$



$$\frac{24.6 - 20.1}{3.0 - 2.0} = 1.5$$

$$RT = 24.6$$

(A) 1.0

(B) 4.5

(C) 1.5

(D) 3.0

S-1

1-14

NCERT ✓

very high .


high →

low .

very low



$$b = 4 \times \frac{4}{3} \pi r^3 \times N_A$$


radius