

College of Engineering  
Mechanical Engineering Department

Second Stage

THERMODYNAMIC I

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CHAPTER THREE – THE WORKING FLUID

LECTURE NO. 2 – THE PERFECT GAS

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# The Perfect Gas

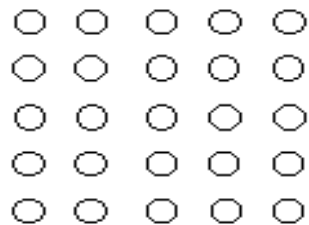
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## THE PERFECT GAS :

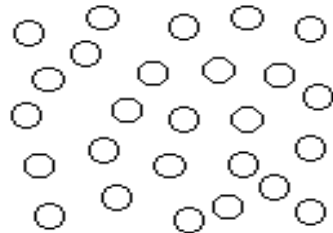
It is a theoretical gas or imaginary gas composed of a set of randomly moving, non-interacting point like particles and they obeys the gas law always.

**OR :** An ideal gas is an imaginary gas that obeys gas law under all conditions

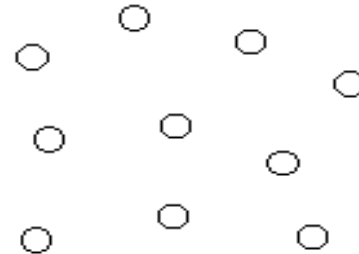
**SOLID**  
atoms are close together, not moving around but are constantly vibrating - they often form a regular pattern



**LIQUID**  
atoms are still close together, but now move around, 'sliding' over each other - there is no regular pattern



**GAS**  
atoms are far apart; they move around freely, but collide with each other and any surfaces they meet



# The Characteristic Equation of State

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## The Characteristic Equation of State :

At temperatures that are considerably in excess of critical temperature of a fluid, and also at very low pressure, the vapour of fluid tends to obey the equation:

$$\frac{pV}{T} = \text{constant} = R$$

### *NOTES :*

- In practice, no gas obeys this law rigidly, but many gases tend towards it.
- Any gas which obeys this law is called a perfect gas,
- This equation is called the **characteristic equation of a state of a perfect gas.**



# The Characteristic Equation of State ..Cont.

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Usually, the characteristic equation is written as

$$pv = RT$$

or for  $m$  kg, occupying  $V$  m<sup>3</sup>

$$pV = mRT$$

Where :

The constant  $R$  is called the *gas constant* (kJ/kg.k).

**Each perfect gas has a different gas constant, See Table 2 Page 2 in Steam Table for value of R**

T : Temperature in Kelvin (K)

p : Pressure in N/m<sup>2</sup>

v : specific volume ( m<sup>3</sup> /kg)



# The Characteristic Equation of State ..Cont.

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The characteristic equation in another form, can be derived by using kilogram-mole as a unit. As per definition of the kilogram-mole, for  $m$  kg of a gas, we have

$$m = nM$$

Where  $n$  = number of moles ,  $M$  : The molecular weight of the gas (kg/mole)

$$pV = nMRT \quad \text{or} \quad MR = \frac{pV}{nT}, \text{ then } MR = R_0$$

As result

$$pV = R_0 nT$$

Where :

$R_0$  : *Universal gas constant ( Fixed value for all gases*

$$R_0 = 8314.3 \text{ Nm/mole K}$$



# The Characteristic Equation of State ..Examples

A vessel of volume  $0.2 \text{ m}^3$  contains nitrogen at  $1.013 \text{ bar}$  and  $15^\circ\text{C}$ . If  $0.2 \text{ kg}$  of nitrogen is now pumped into the vessel, calculate the new pressure when the vessel has returned to its initial temperature. The molar mass of nitrogen is  $28 \text{ kg/kmol}$ , and it may be assumed to be a perfect gas.

$$\text{Specific gas constant, } R = \frac{\bar{R}}{\bar{m}} = \frac{8314.5}{28} = 296.95 \text{ N m/kg K}$$

From equation (2.6), for the initial conditions

$$p_1 V_1 = m_1 R T_1$$

therefore

$$m_1 = \frac{p_1 V_1}{R T_1} = \frac{1.013 \times 10^5 \times 0.2}{296.95 \times 288} = 0.237 \text{ kg}$$

where  $T_1 = 15 + 273 = 288 \text{ K}$

The mass of nitrogen added is  $0.2 \text{ kg}$ , hence  $m_2 = 0.2 + 0.237 = 0.437 \text{ kg}$ .  
for the final conditions

$$p_2 V_2 = m_2 R T_2$$

but  $V_2 = V_1$  and  $T_2 = T_1$ , therefore

$$p_2 = \frac{m_2 R T_2}{V_2} = \frac{0.437 \times 296.95 \times 288}{10^5 \times 0.2}$$

i.e.  $p_2 = 1.87 \text{ bar}$



# Specific Heats

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## Specific Heats

The **specific heat** of a solid or liquid is usually defined as the heat required to raise unit mass through one degree temperature rise.

For small quantities, we have :  $dQ = mcdT$

where

$m$  = mass,

$c$  = specific heat

$dT$  = temperature rise.



# Specific Heats..Cont.

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We have two specific heats for gases are defined.

- *Specific heat at constant volume,  $c_v$*  and,
- *Specific heat at constant pressure,  $c_p$*

The heat flow between two temperature can defined as :

$$dQ = m c_p dT \quad \text{For a reversible non-flow process at } \textit{constant pressure}$$

and,

$$dQ = m c_v dT \quad \text{For a reversible non-flow process at } \textit{constant volume}$$





# Specific Heats..Cont.

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The values of  $c_p$  and  $c_v$ , for a perfect gas, are constant for any one gas at all pressures and temperatures , we have

**Flow of heat in a reversible constant pressure process,  $Q = mc_p (T_2 - T_1)$**

**Flow of heat in a reversible constant volume process,  $Q = mc_v (T_2 - T_1)$**

In case of real gases,  $c_p$  and  $c_v$  vary with temperature, but a suitable average value may be used for most practical purposes.



# Joule's Law



## Joule's Law

### ( Definition of Internal energy for perfect gas ) :

The internal energy of a perfect gas is a function of the absolute temperature only. i.e.,  
 $u = f(T)$

Specific Internal energy,  $u = c_v T$  for a perfect gas

or For mass  $m$ , of a perfect gas

Internal energy,  $U = mc_v T$

For a perfect gas, in any process between states 1 and 2, we have

Gain in internal energy,  $U_2 - U_1 = mc_v (T_2 - T_1)$



# Relationship Between Two Specific Heats

## Relationship Between Two Specific Heats

Consider a perfect gas being heated at constant pressure from  $T_1$  to  $T_2$ . According to non-flow equation,  $Q = (U_2 - U_1) + W$

Also for a perfect gas,  $U_2 - U_1 = mc_v (T_2 - T_1)$  and  $Q = mc_v (T_2 - T_1) + W$

In a constant pressure process, the work done by the fluid,  $W = p (V_2 - V_1) = mR (T_2 - T_1)$  ,  $pV = mRT$  ,

In this case On substituting

$Q = mc_v (T_2 - T_1) + mR (T_2 - T_1) = m(c_v + R) (T_2 - T_1)$  But for a constant pressure process,  $Q = mc_p (T_2 - T_1)$  ,

By equating the two expressions, we have

$$m(c_v + R)(T_2 - T_1) = mc_p(T_2 - T_1)$$

$$\therefore c_v + R = c_p \quad \text{or} \quad c_p - c_v = R \quad \text{Dividing both sides by } c_v, \text{ we get } \frac{c_p}{c_v} - 1 = \frac{R}{c_v} ,$$

$$\gamma = \frac{c_p}{c_v} \quad \text{Ratio of Specific Heats}$$

The ratio of specific heat at constant pressure to the specific heat at constant volume is given the symbol  $\gamma$  (gamma). Then

$$c_v = \frac{R}{\gamma - 1} \quad \text{and} \quad c_p = \frac{\gamma R}{\gamma - 1}$$



# Enthalpy of Perfect Gas

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## Enthalpy

One of the fundamental quantities which occur invariably in thermodynamics is the sum of internal energy ( $u$ ) and pressure volume product ( $p\nu$ ). This sum is called **Enthalpy (h)**.

$$\text{i.e., } h = u + p\nu$$

The total enthalpy of mass,  $m$ , of a fluid can be  $H = U + pV$ , where  $H = mh$ .

For a **perfect gas**,  $h = u + p\nu = cvT + RT$       Where  $p\nu = RT$

$= (cv + R)T = cpT$       Where  $cp = cv + R$

$$h = cpT \text{ and } H = mcpT$$

