

Properties of Substances

Presentation Outline

- Overview
- Property Diagrams
- Property Tables

Overview

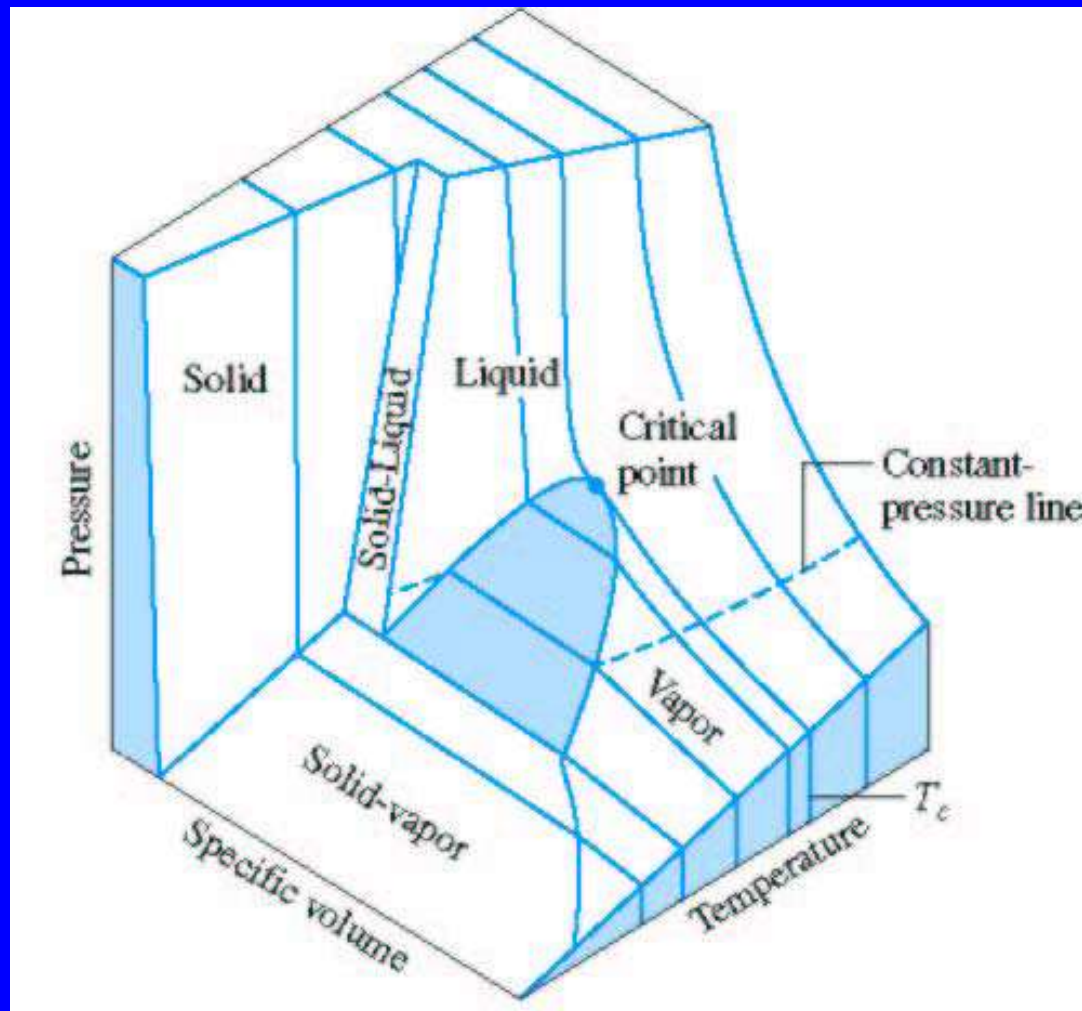
- A *pure substance* is one of uniform and invariable chemical composition.
- For present analysis, a simple substance is taken as one for which if the values of two intensive properties are known, all the other properties can be found. E.g. $u = f(T, p)$

Overview contd..

- A material can exist in the
 - Solid phase
 - Liquid phase
 - Vapour (or Gaseous) phase
 - A mixture of the above phases in equilibrium (e.g. melting, vaporization or sublimation)
- Through experiments it is known that temperature and specific volume can be considered as independent variables and pressure can be determined as a function of the two.

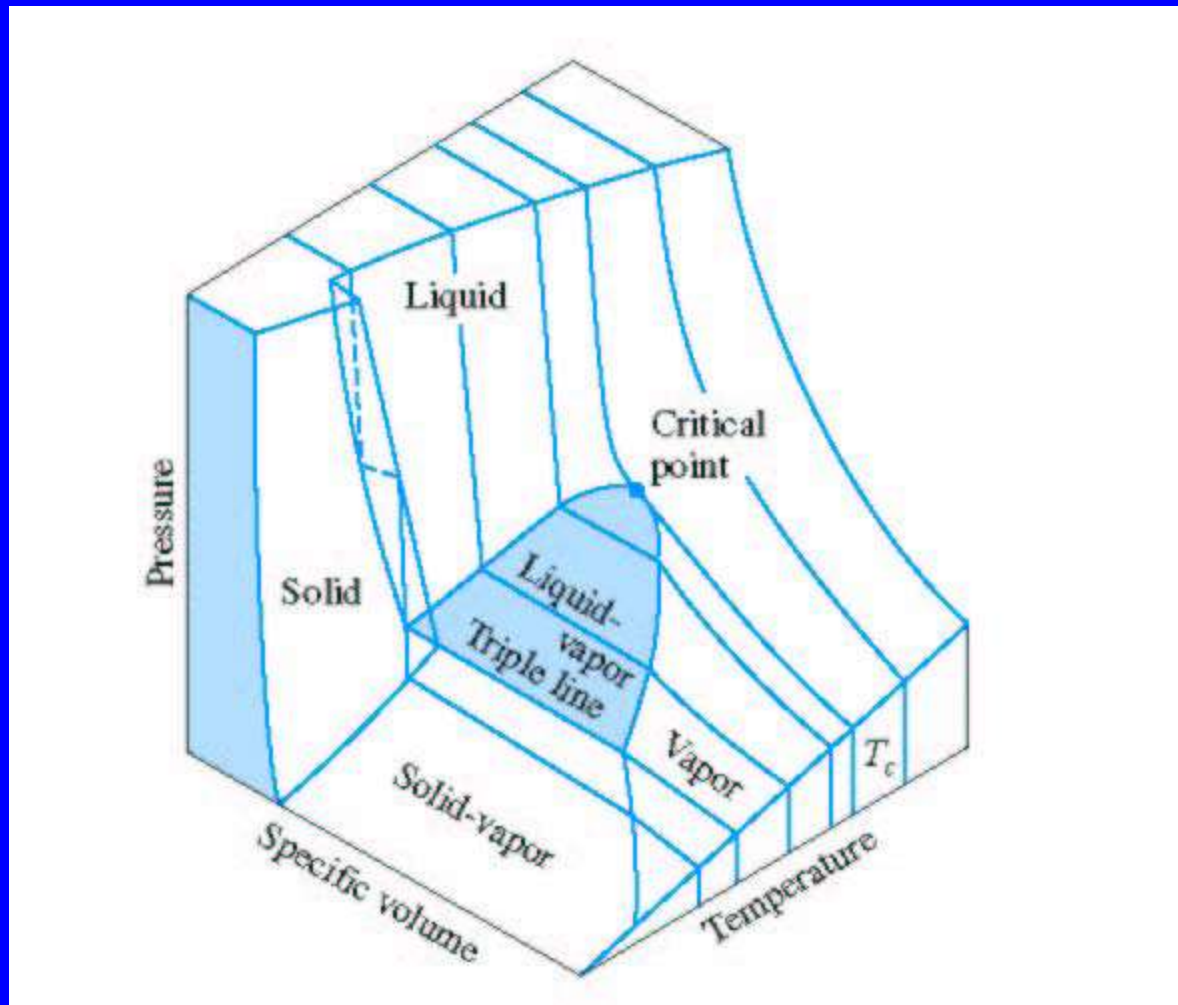
$$p = f(T, v)$$

p-v-T plot of a common metal



- This creates a p-v-T surface on the plot

p-v-T plot of water



- Note the step increase in specific volume (step decrease in density) when transferring from liquid to solid

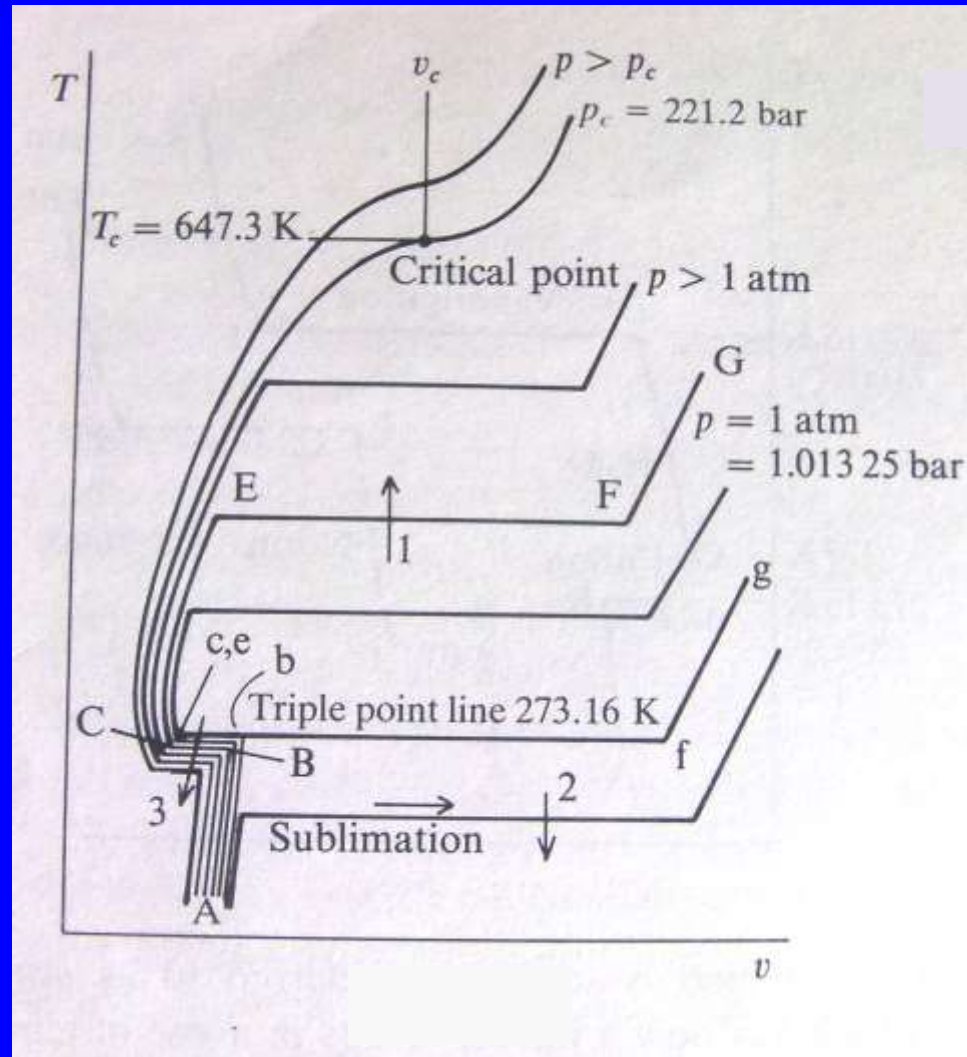
Property Diagrams

- 3-D surface plots are not very useful and rarely used.
- 2-D projections are more instructive from analytical point of view.
- Different standard property diagrams are used for obtaining relationships among different properties of substances.

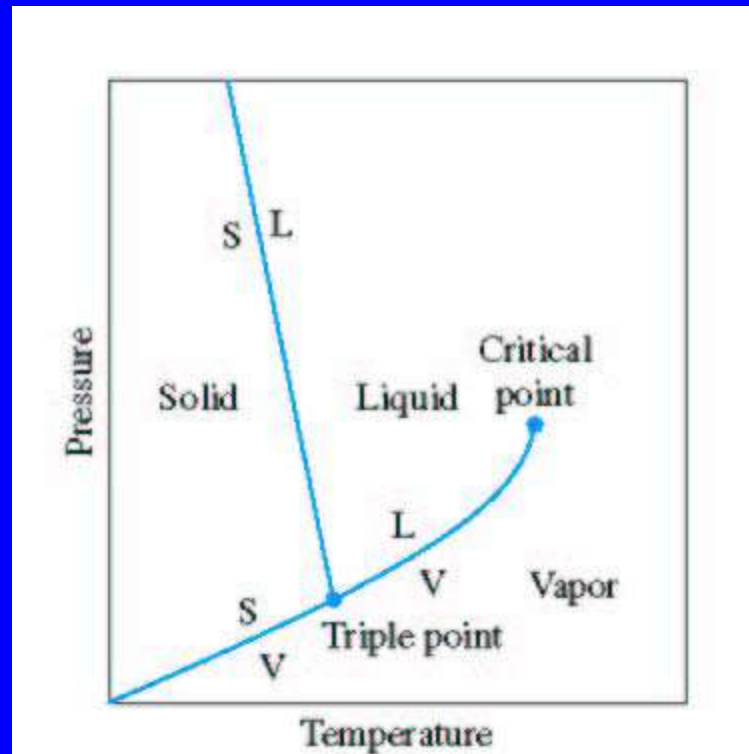
Property Diagrams contd..

- Common 2-D property diagrams include
 - Temperature vs. Specific Volume diagram (T-v diagram)
 - Pressure vs. Specific Volume diagram (p-v diagram)
 - Temperature vs. Specific Entropy diagram (T-s diagram)
 - Specific Enthalpy vs. Specific Entropy diagram (h-s diagram)
 - Pressure vs. Enthalpy diagram (p-h diagram)

T-v diagram for water

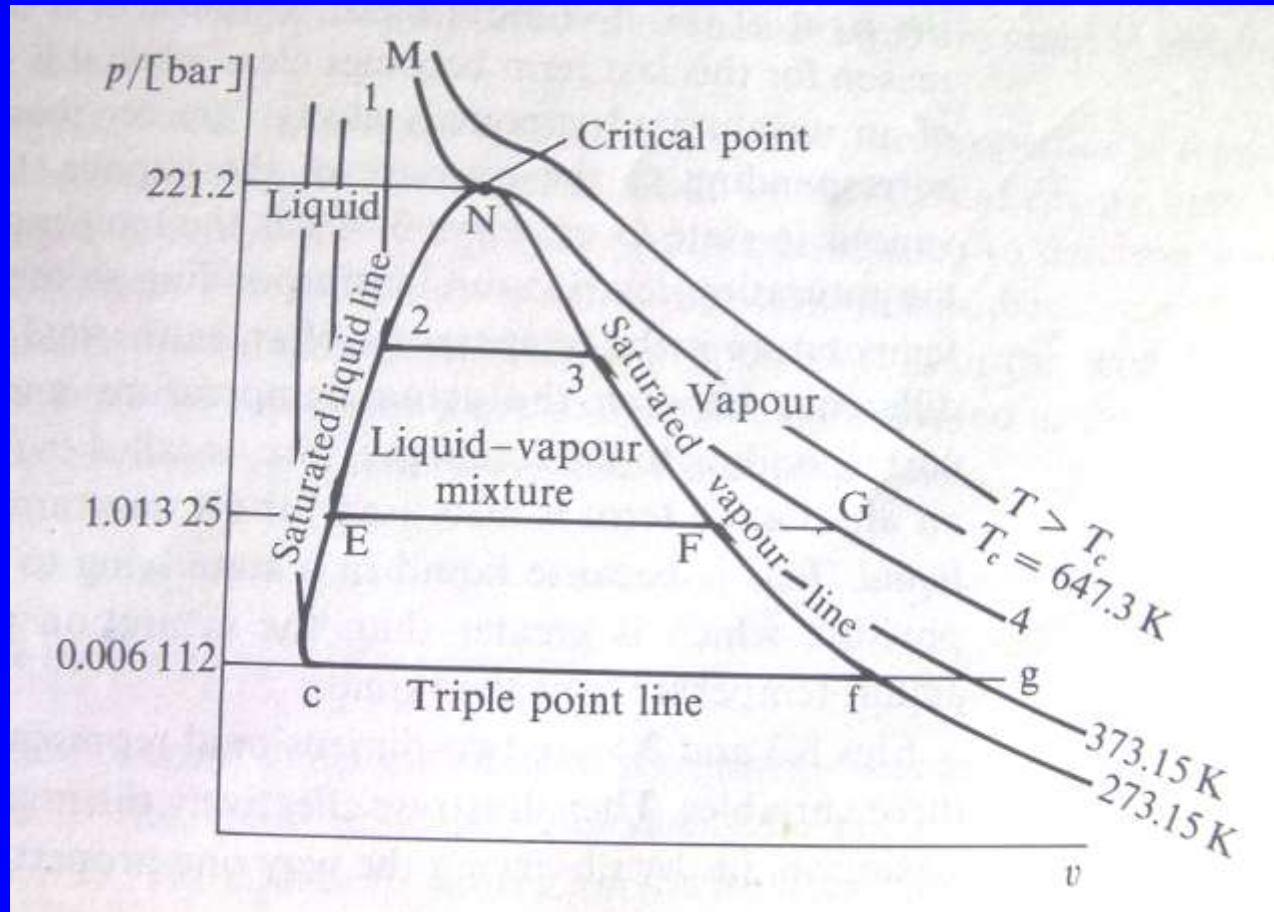


p-T diagram for water



- Only single-phase regions observed
- Two-phase regions appear as lines (edge view)
- Triple point is represented on the edge view of the triple line where all three phases co-exist in equilibrium.

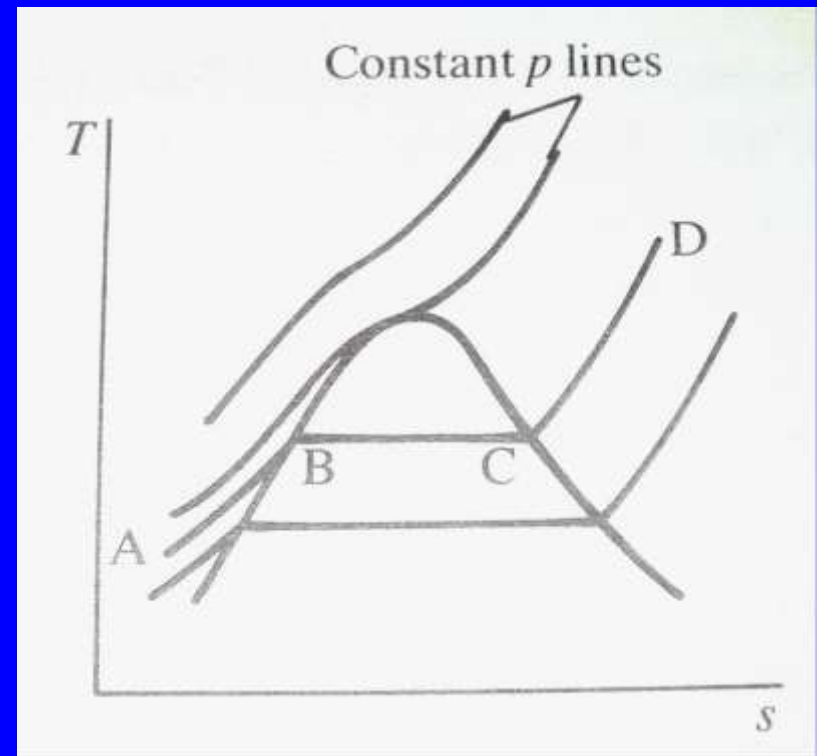
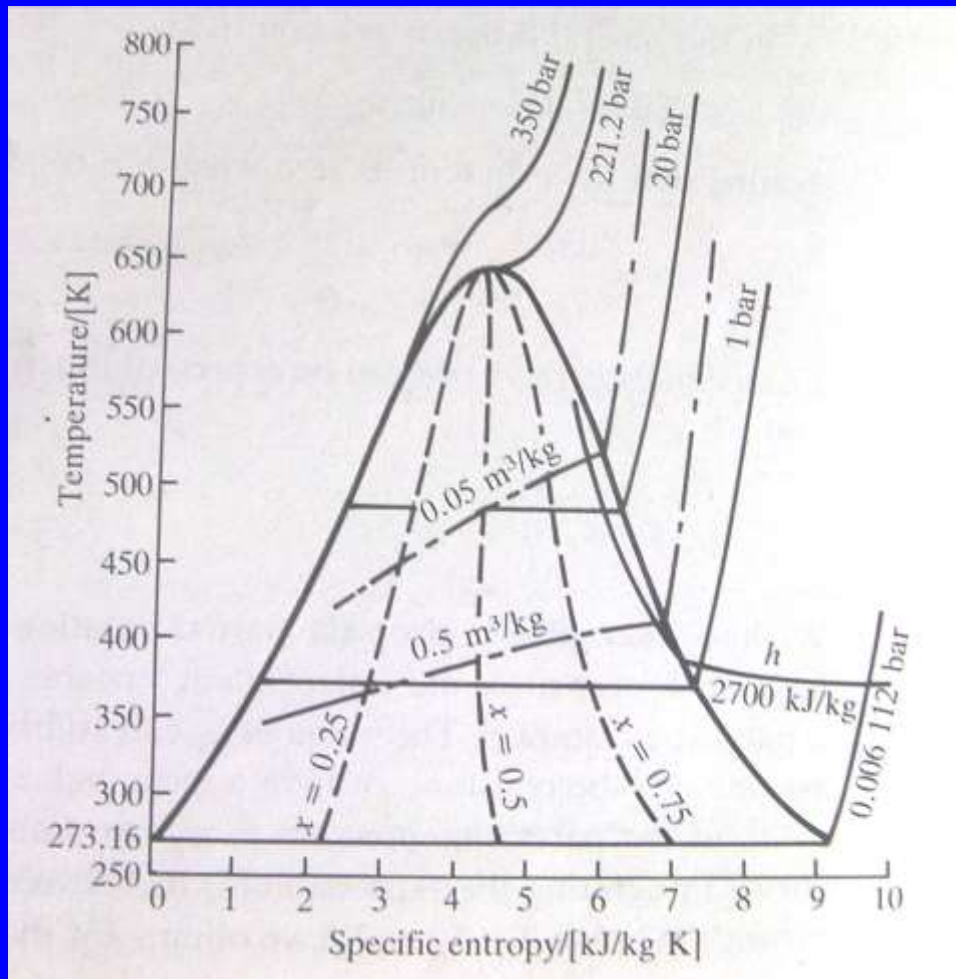
p-v diagram for water



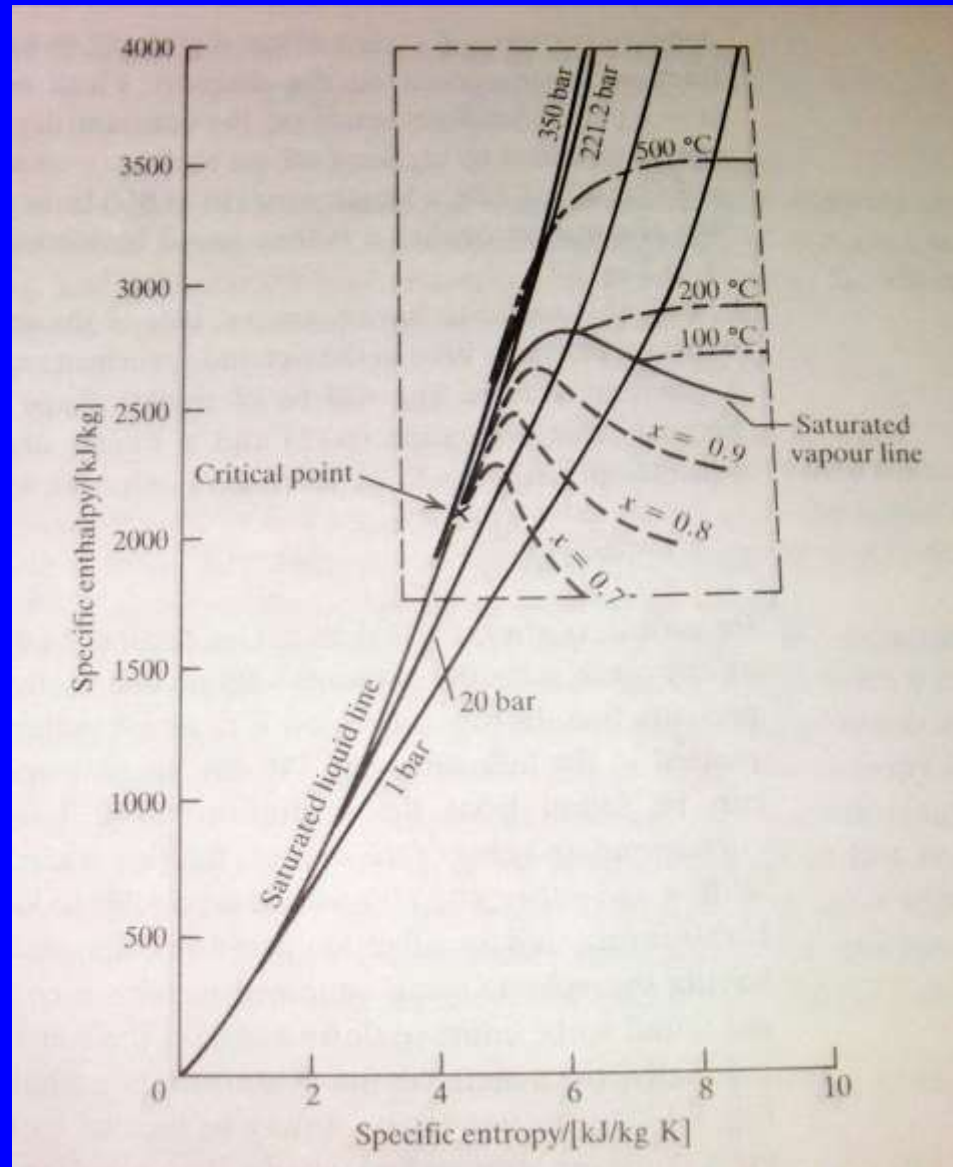
p-v diagram for water contd..

- The dome-shaped two phase region is comprised of liquid and vapour, is called the saturation dome.
- Constant temperature lines shown are called isotherms.
- Above the critical point isobars and isotherms do not pass through the liquid-vapour region.
- The critical point defines the maximum pressure. i.e. Above the critical pressure liquid and vapour cannot co-exist in equilibrium.
- In single-phase regions, the state is fixed by any two of the properties p , v and T that are independent variables.
- In two-phase regions, p and T are not independent and the state cannot be fixed by p and T but can be explained by p and v or T and v .

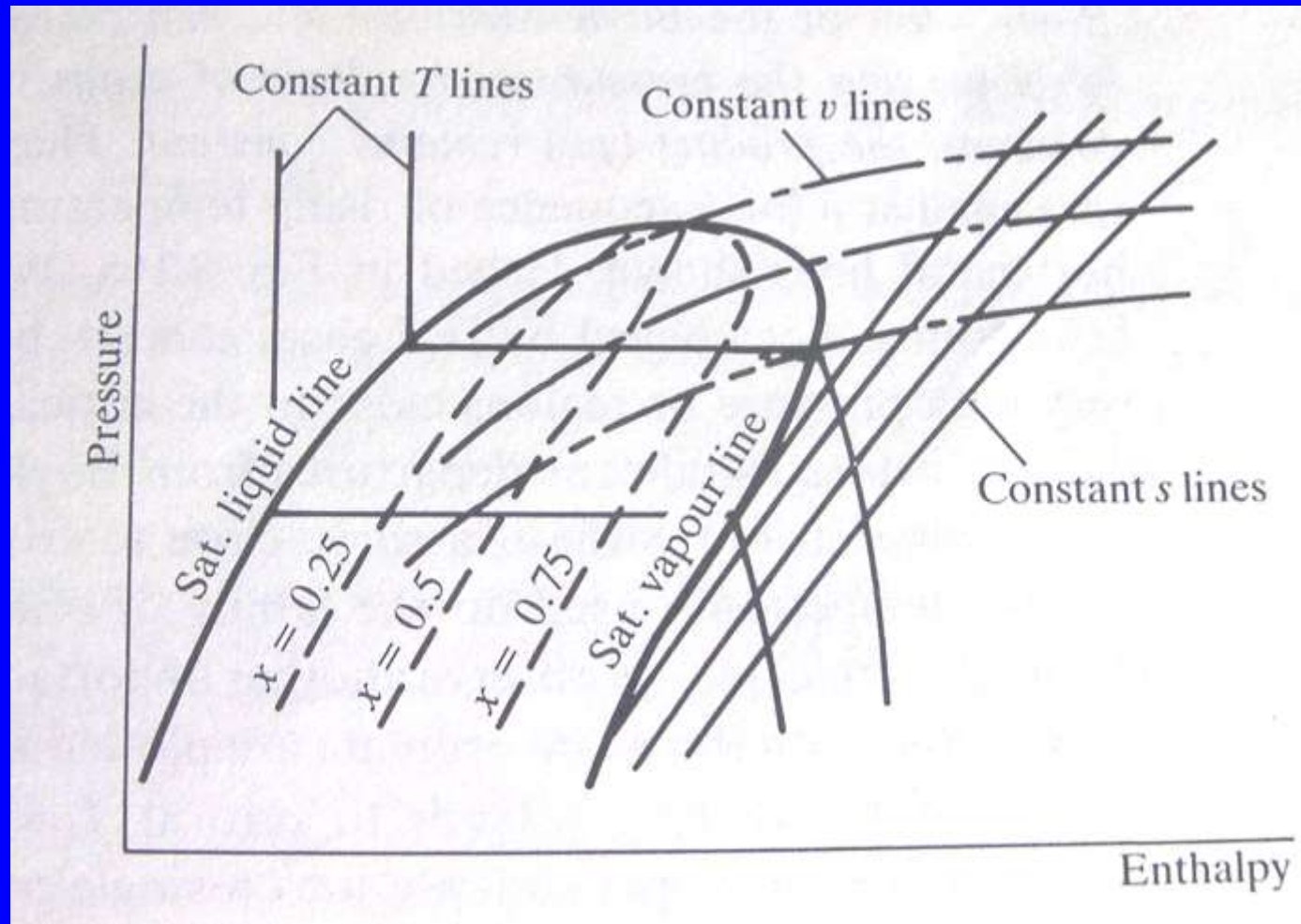
T- s diagram for water



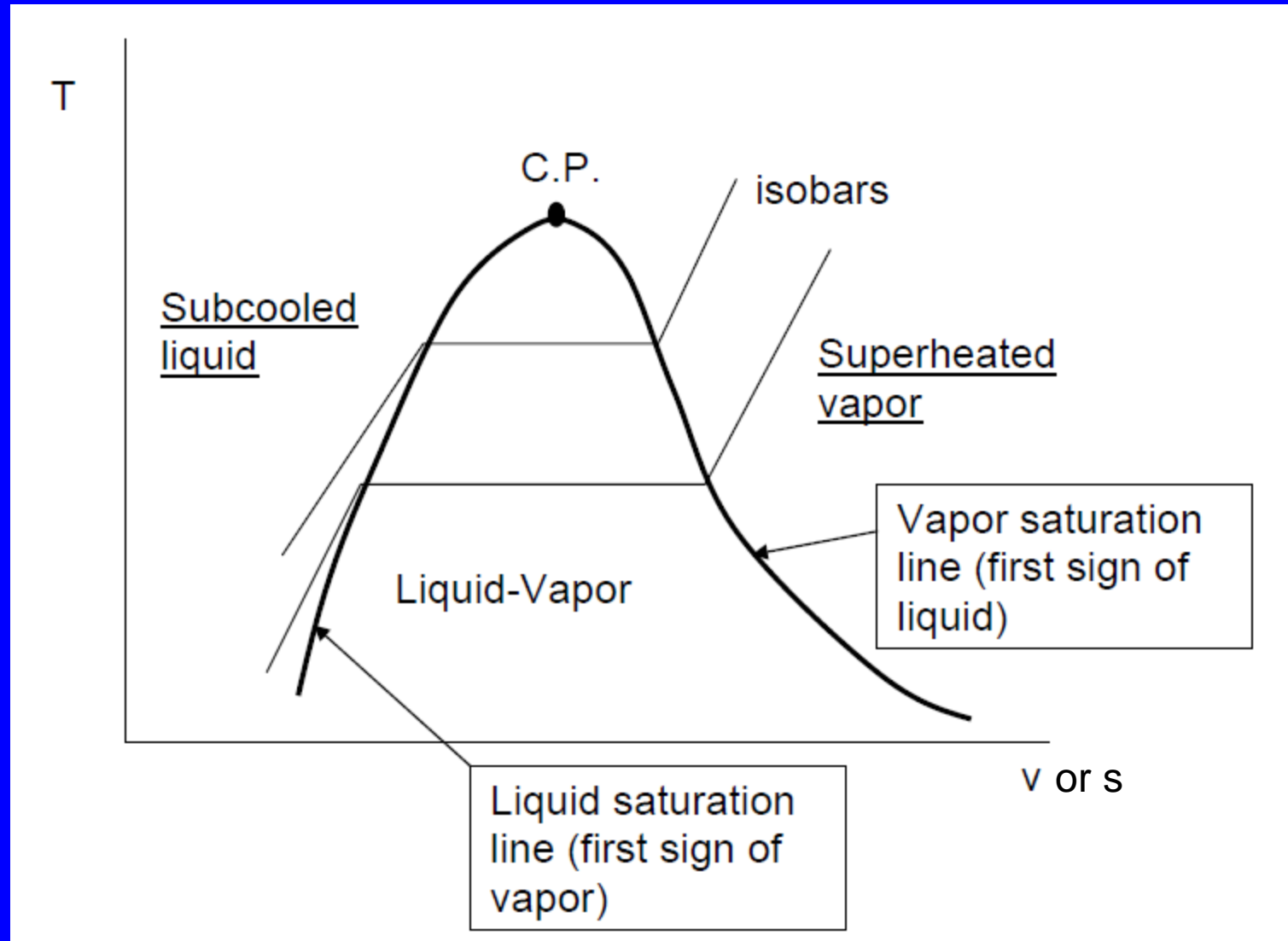
h-s diagram for water



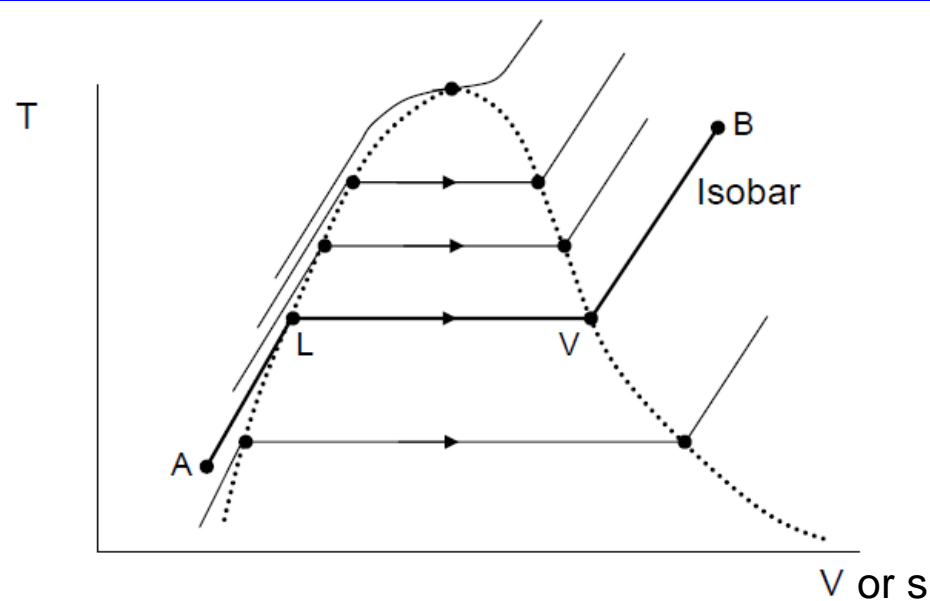
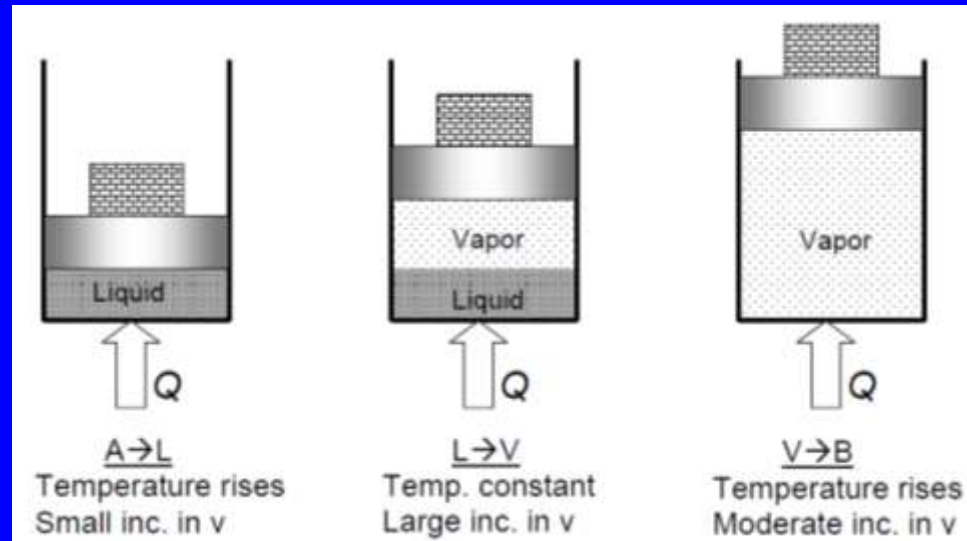
p-h diagram for water



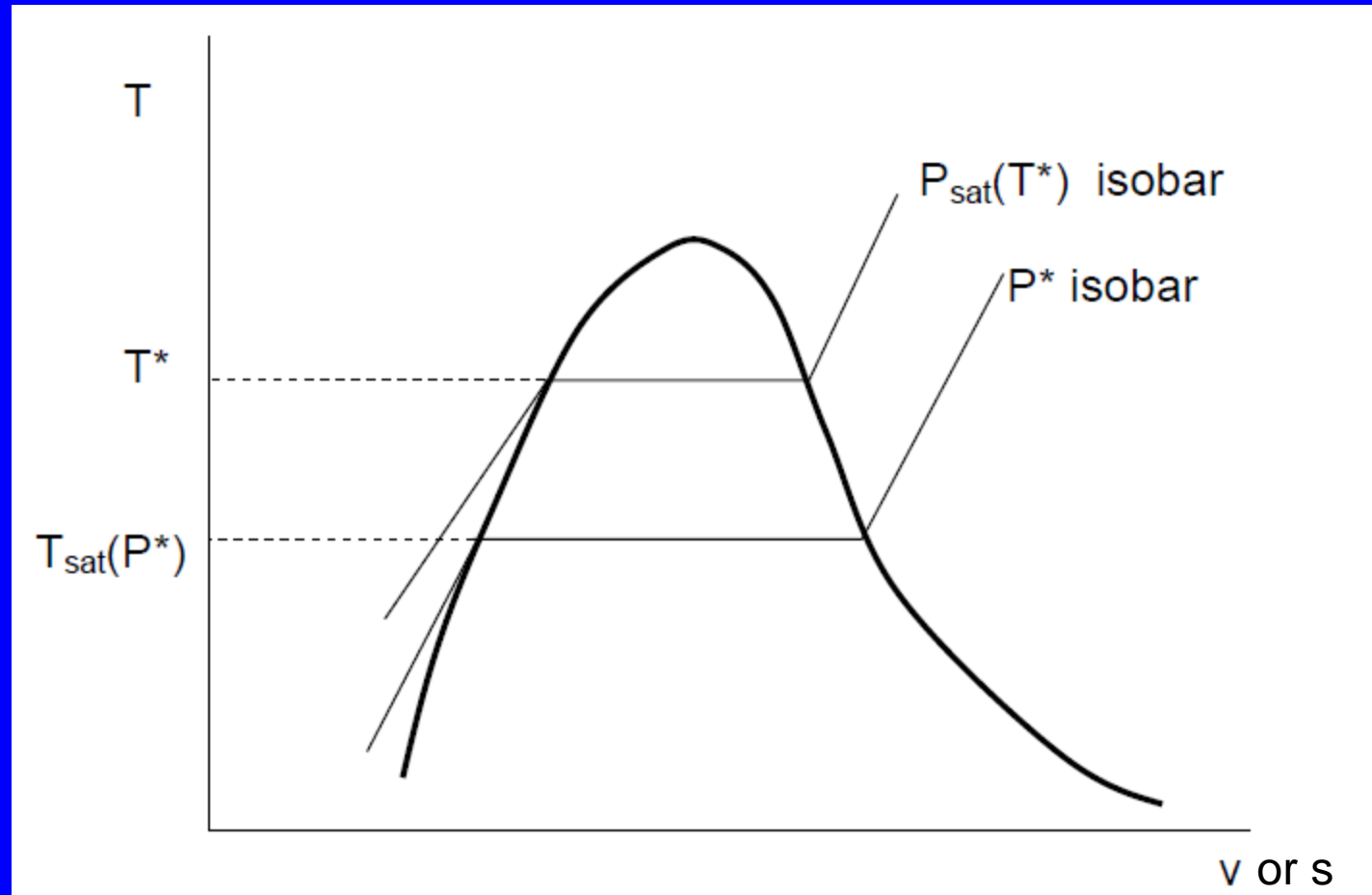
Characteristics of Saturation Dome



Consider the following constant pressure heating process



Saturation Pressure & Temperature

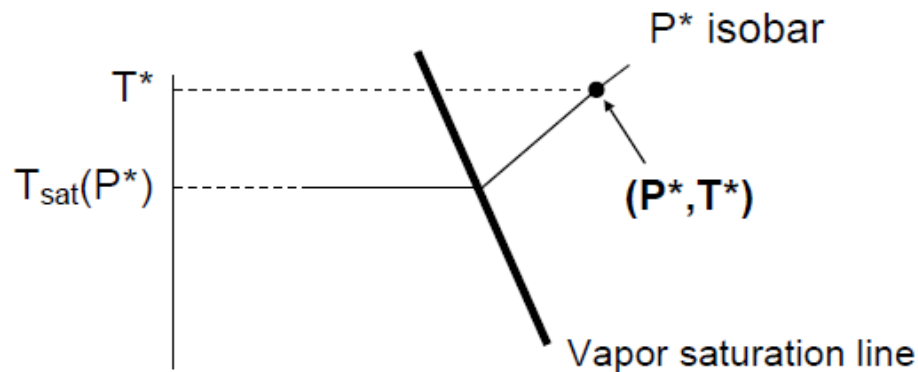


Saturation Pressure & Temperature

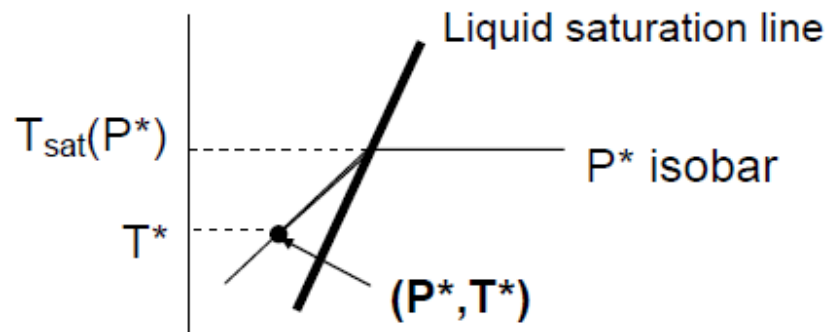
- **Saturation Temperature:** Temperature of a liquid-vapour mixture in equilibrium at a pressure P^* , denoted as $T_{\text{sat}}(P^*)$. This is the temperature where the P^* isobar crosses vapour saturation line.
- **Saturation Pressure:** Pressure of a liquid-vapour mixture in equilibrium at a temperature T^* , denoted as $P_{\text{sat}}(T^*)$. This is the pressure at which the T^* isotherm crosses the liquid saturation line.

Saturation Pressure & Temperature contd..

Superheated Vapor: $T^* > T_{\text{sat}}(P^*)$



Subcooled liquid: $T^* < T_{\text{sat}}(P^*)$



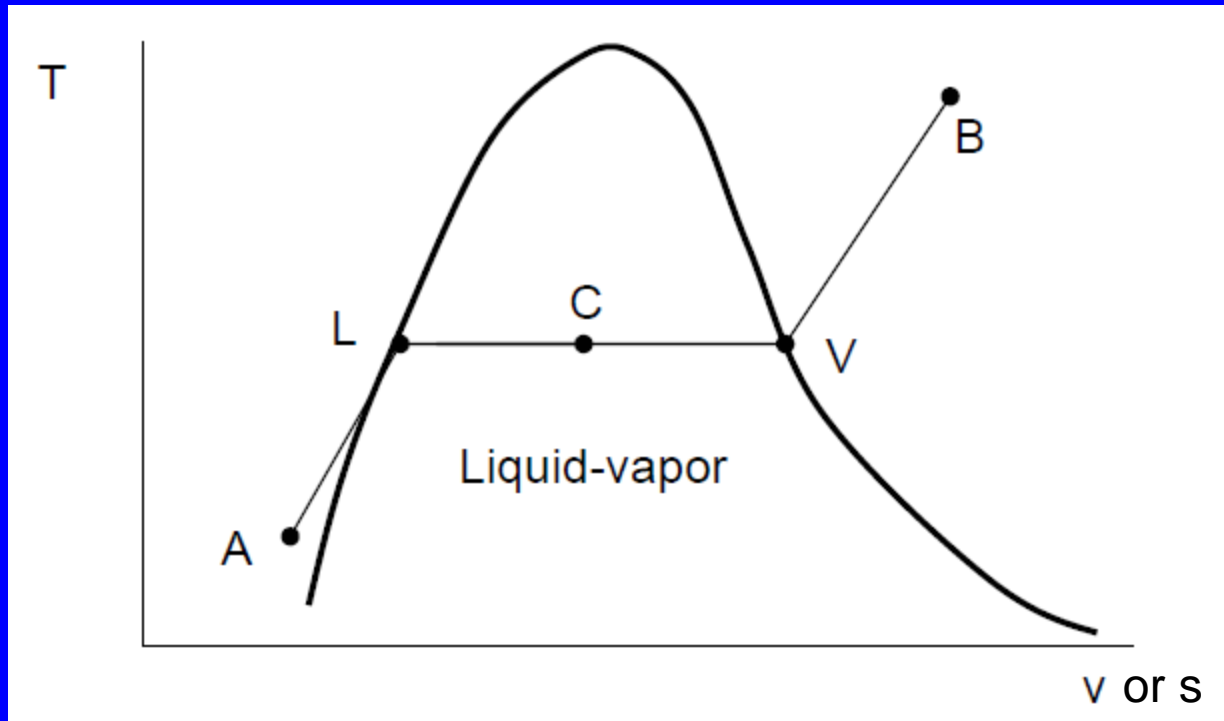
Dryness Fraction (x)

- This gives an indication of the quality and characteristics of steam being utilized.
- It is defined as the ratio of the mass of vapour (m_v) to the total mass of a two-phase liquid-vapour mixture (m).
- Mass of liquid = m_l

Dryness Fraction

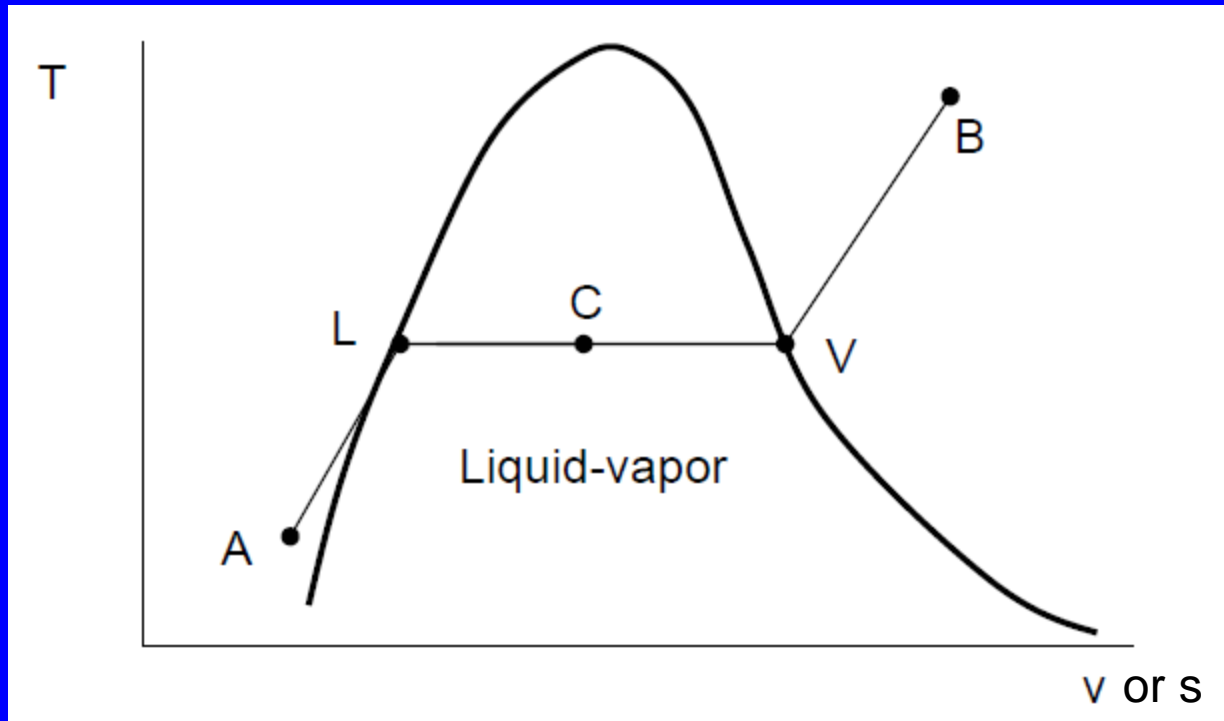
$$x = \frac{m_v}{m} = \frac{m_v}{m_v + m_l}$$

Dryness Fraction contd..



- At point L, $x = 0$
- At point V, $x = 1$
- Between L and V, $0 < x < 1$
- Dryness Fraction may also be presented as a percentage ($0 \leq x \leq 100$)

Liquid – Vapour Properties



- Properties at point L are denoted by subscript f. (E.g. u_f , v_f , s_f , h_f etc.)
- Properties at point V are denoted by subscript g. (E.g. u_g , v_g , s_g , h_g etc.)

Liquid –Vapour Properties contd..

- Let us consider an intermediate point C and its corresponding properties are u , v , s and h .
- Then the following expressions will provide the properties of the two-phase fluid at C.

1. Specific Internal Energy

$$u = (1 - x)u_f + xu_g \quad \text{or} \quad u = u_f + x(u_g - u_f)$$

$$\text{If } u_{fg} = u_g - u_f \quad \text{then} \quad u = u_f + xu_{fg}$$

Liquid – Vapour Properties contd..

2. Specific Volume

$$v = (1 - x)v_f + xv_g$$

However v_f is very small compared with v_g .

Hence
$$v = xv_g$$

3. Specific Entropy

$$s = (1 - x)s_f + xs_g \quad \text{or} \quad s = s_f + x(s_g - s_f)$$

If $s_{fg} = s_g - s_f$ then
$$s = s_f + xs_{fg}$$

Liquid – Vapour Properties contd..

4. Specific Enthalpy

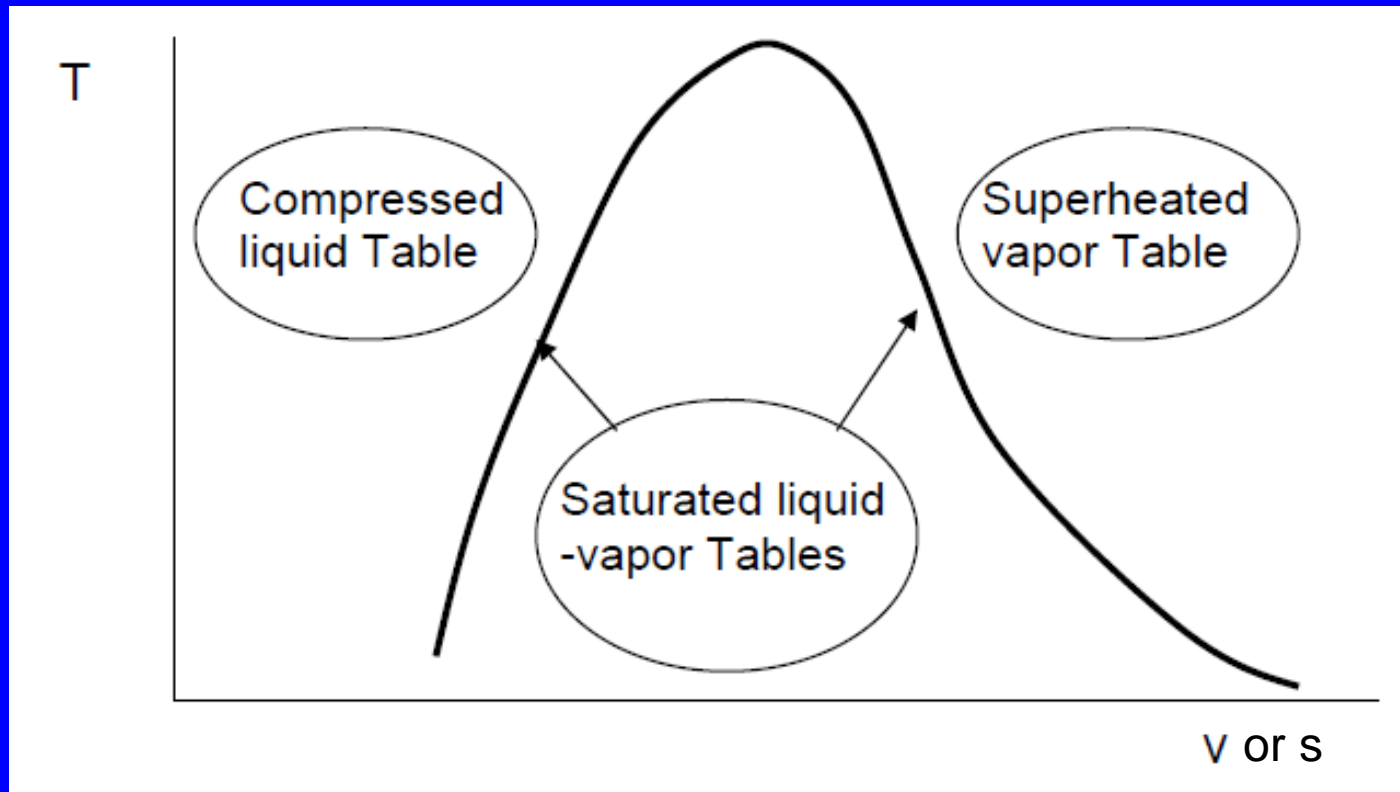
$$h = (1 - x)h_f + xh_g \quad \text{or} \quad h = h_f + x(h_g - h_f)$$

$$\text{If } h_{fg} = h_g - h_f \quad \text{then} \quad h = h_f + xh_{fg}$$

- Hence for any property α at C is given by

$$\alpha = (1 - x)\alpha_f + x\alpha_g \quad \text{or} \quad \alpha = \alpha_f + x\alpha_{fg}$$

Use of Property Tables



- Appropriate Property Tables are used to define the relationships among different properties of fluids.

Properties of Superheated Steam

- For steam in the superheated region, temperature and pressure are independent properties.
- When the temperature and pressure are given for superheated steam then the state is defined and all the other properties can be found.