

Some Consequences of T_{sat} and P_{sat} Dependence

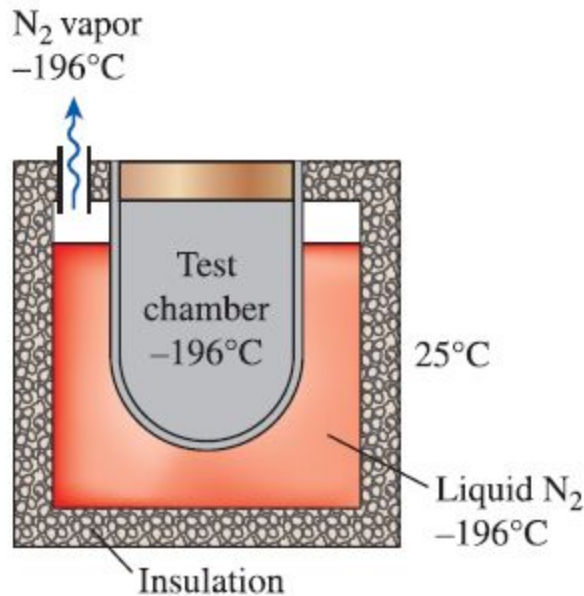
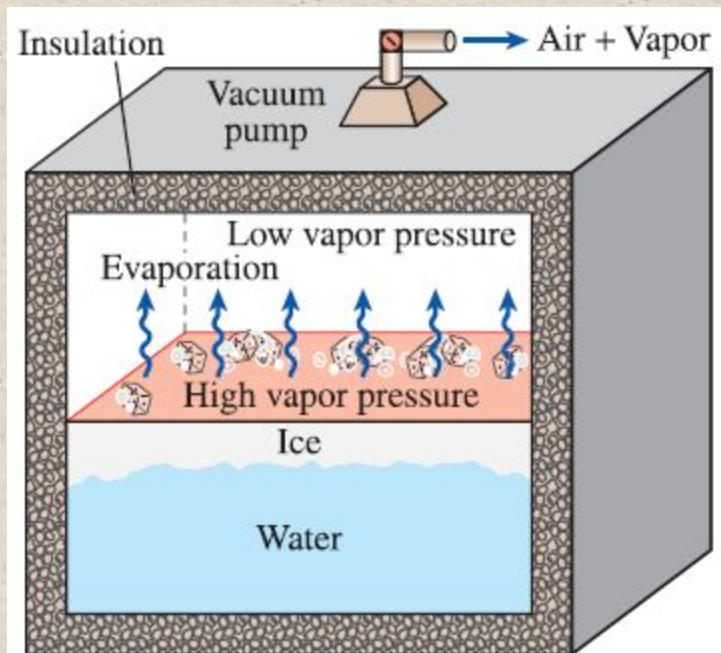
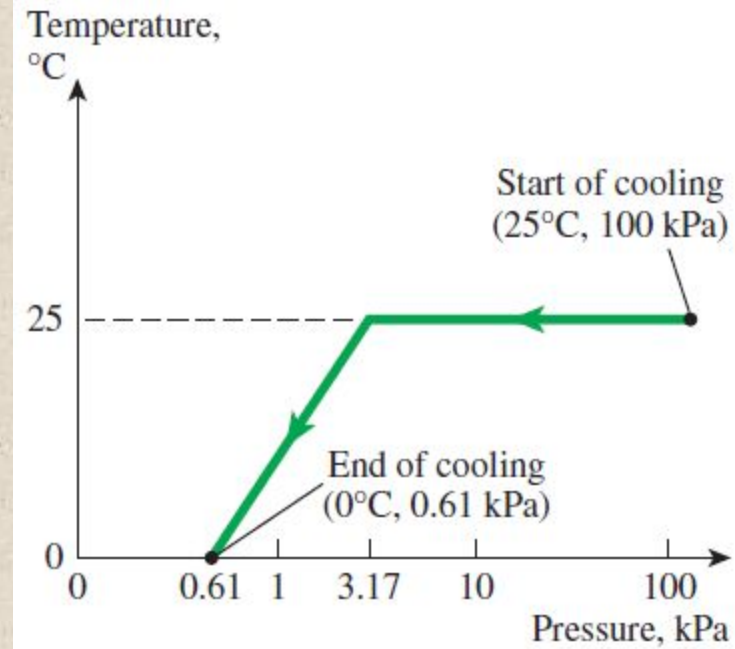


FIGURE 3-12

The temperature of liquid nitrogen exposed to the atmosphere remains constant at -196°C , and thus it maintains the test chamber at -196°C .

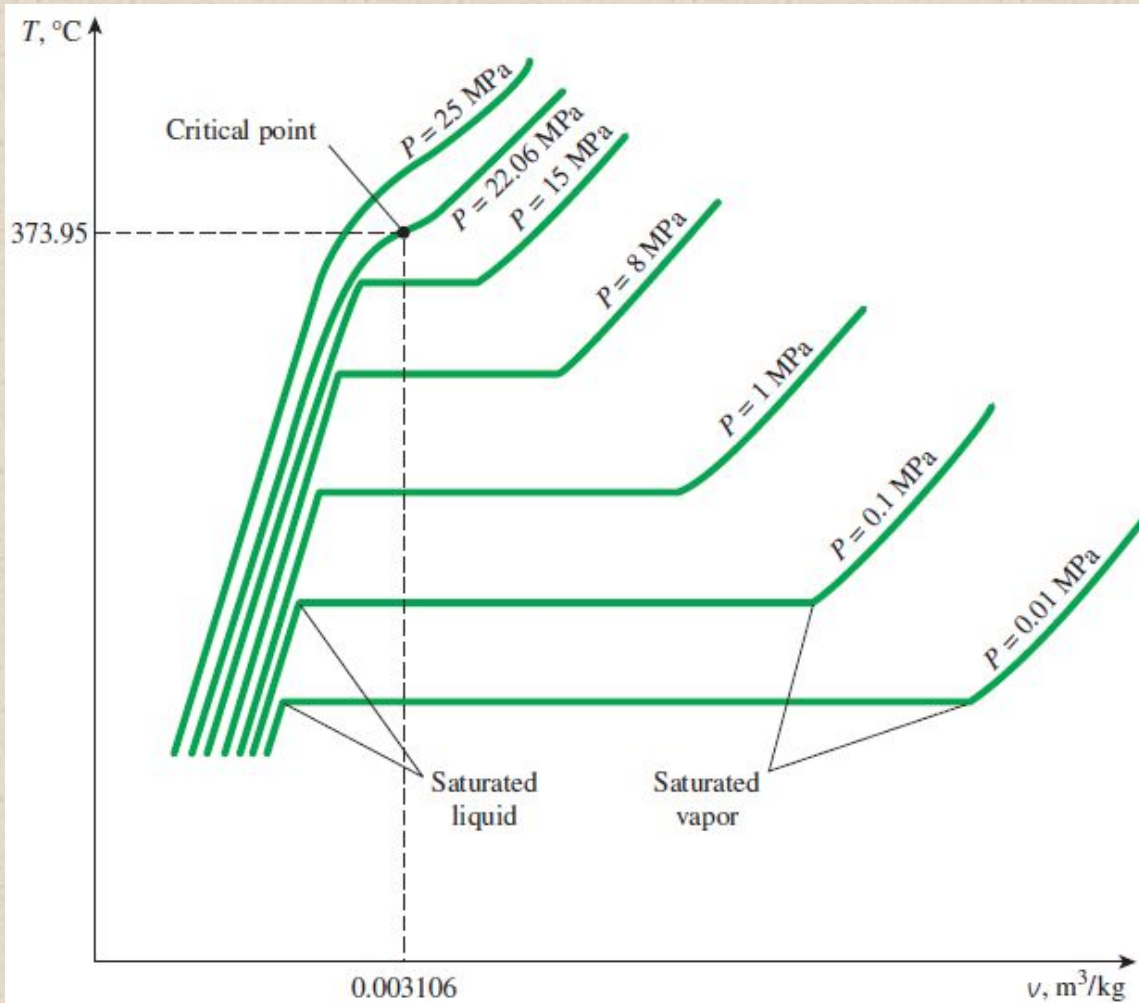
The variation of the temperature of fruits and vegetables with pressure during vacuum cooling from 25°C to 0°C .



In 1775, ice was made by evacuating the air space in a water tank.

PROPERTY DIAGRAMS FOR PHASE-CHANGE PROCESSES

The variations of properties during phase-change processes are best studied and understood with the help of property diagrams such as the T - v , P - v , and P - T diagrams for pure substances.



T - v diagram of constant-pressure phase-change processes of a pure substance at various pressures (numerical values are for water).

- saturated liquid line
- saturated vapor line
- compressed liquid region
- superheated vapor region
- saturated liquid–vapor mixture region (wet region)

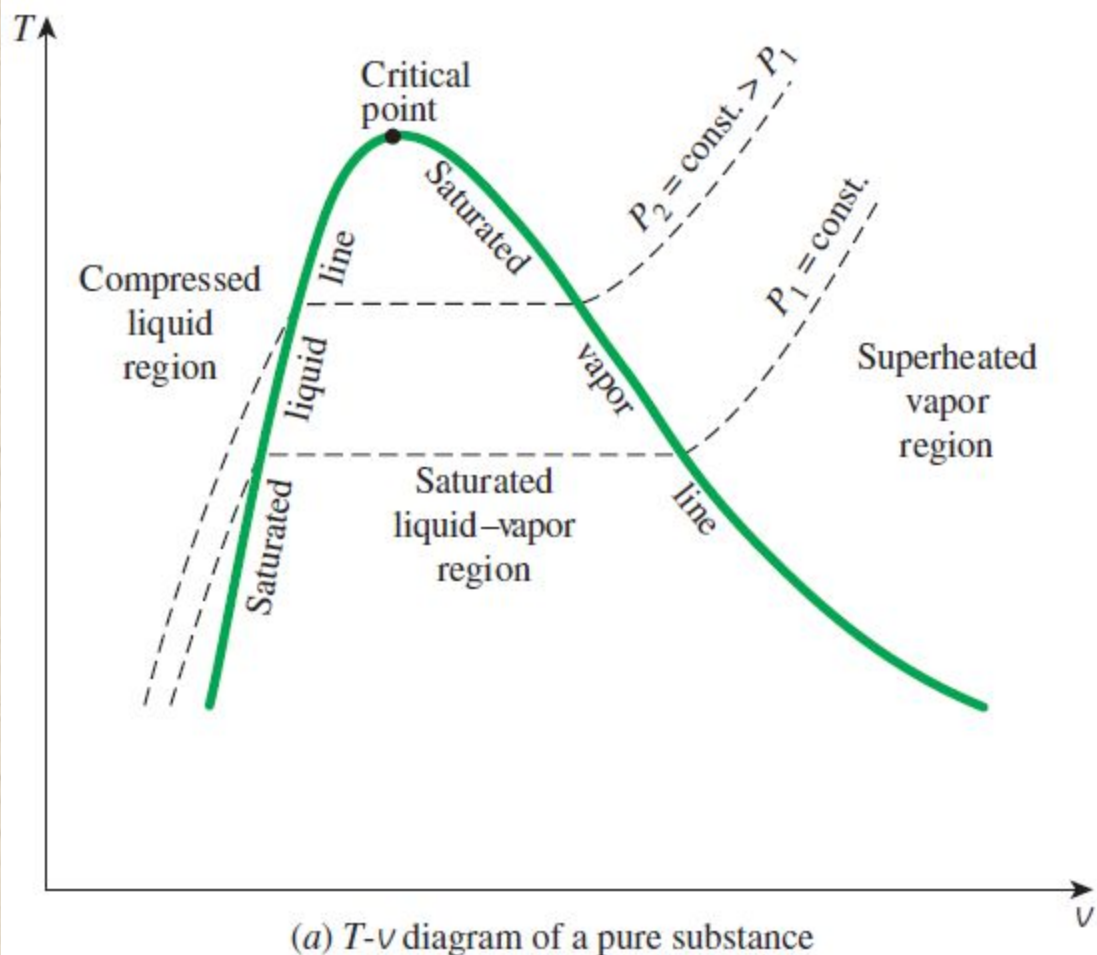
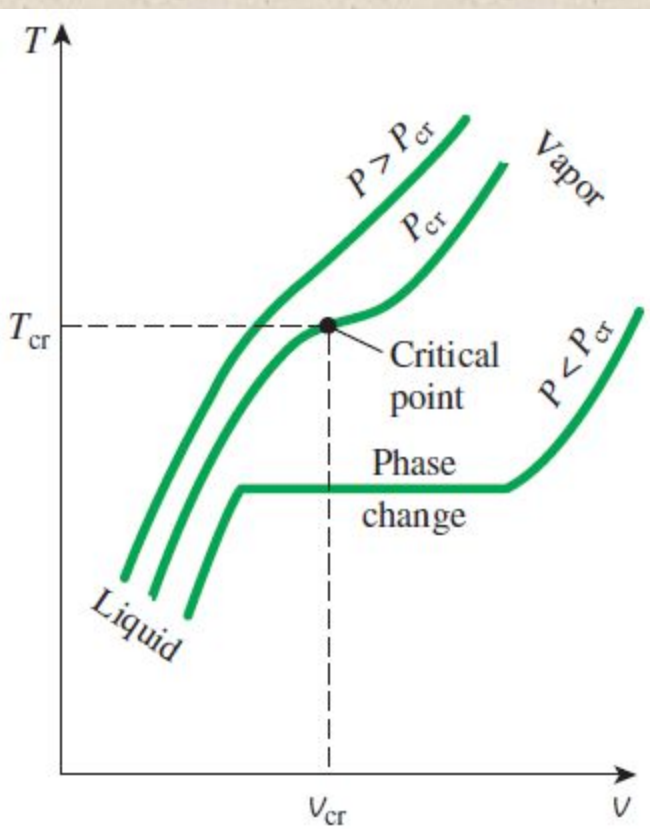


FIGURE 3–17

Property diagrams of a pure substance.



At supercritical pressures ($P > P_{cr}$), there is no distinct phase-change (boiling) process.

Critical point: The point at which the saturated liquid and saturated vapor states are identical.

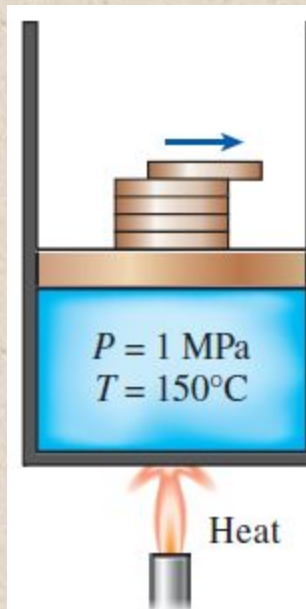
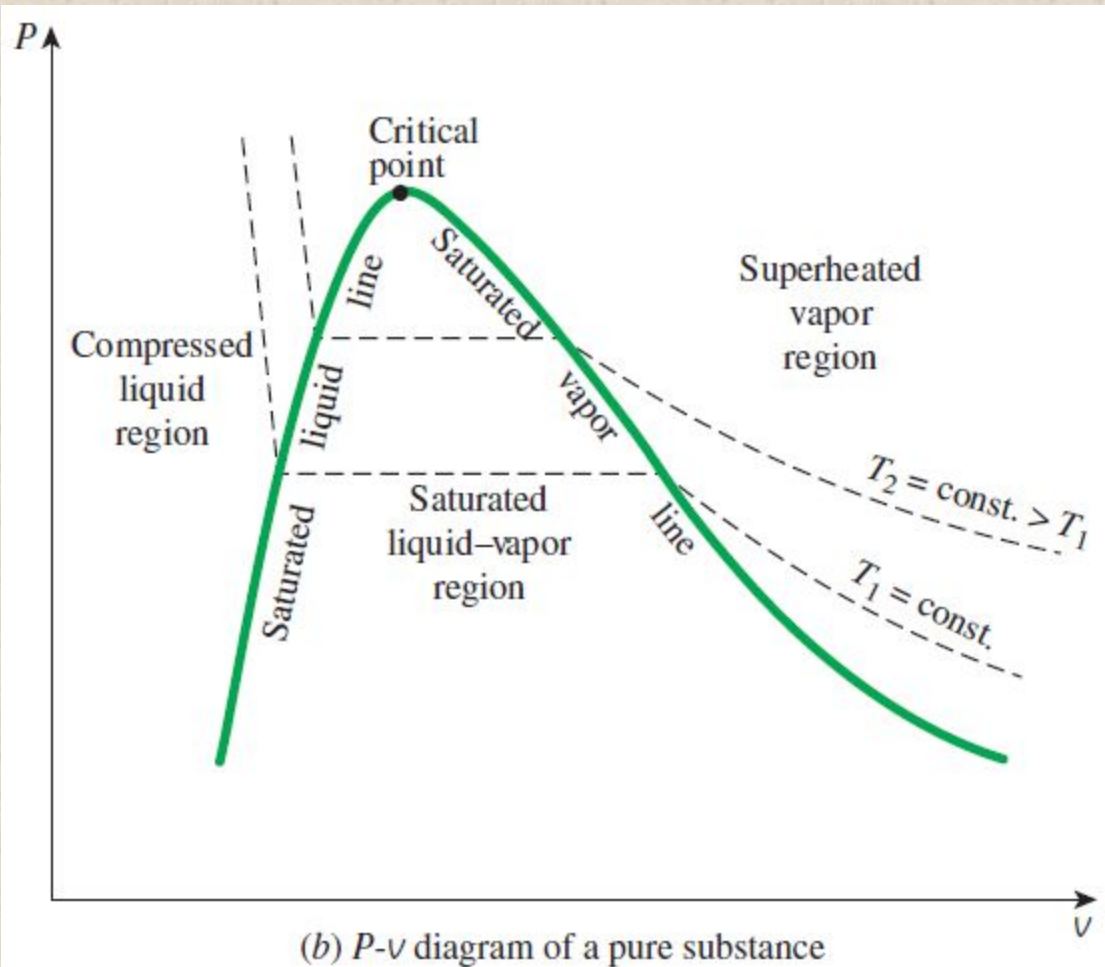


FIGURE 3-18

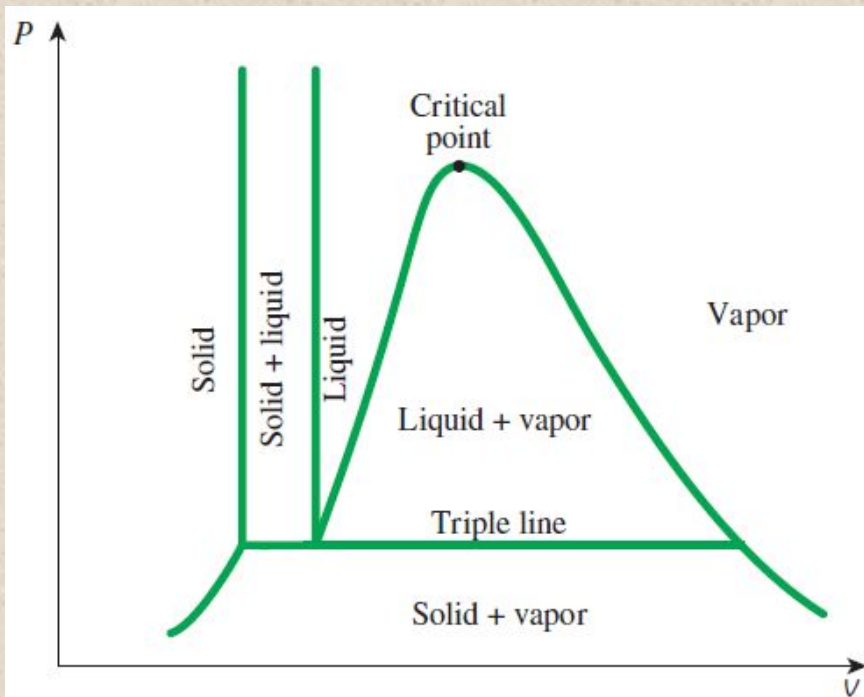
The pressure in a piston-cylinder device can be reduced by reducing the weight of the piston.

Extending the Diagrams to Include the Solid Phase

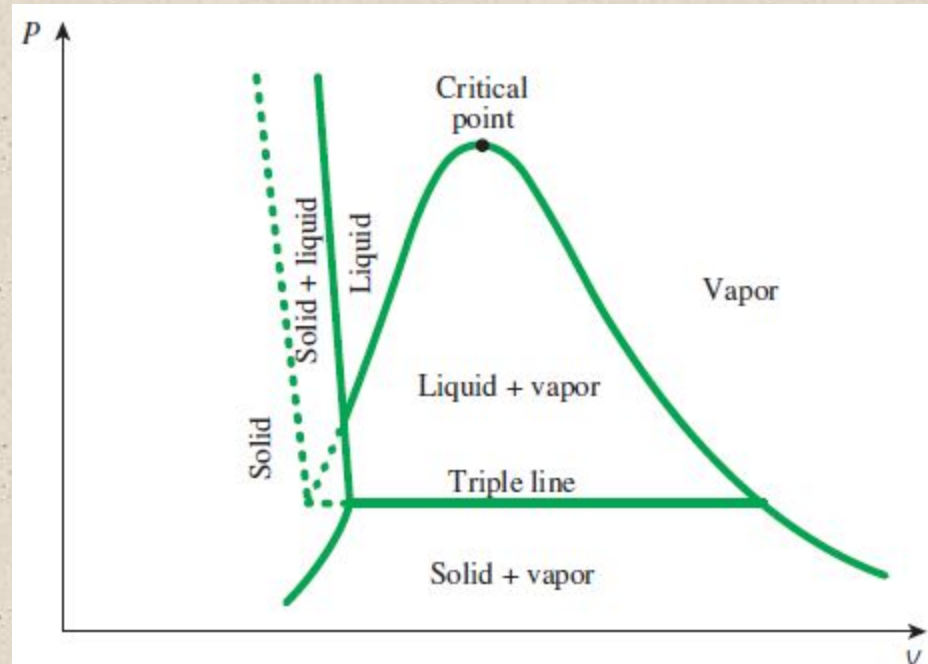
For water,
 $T_{tp} = 0.01^{\circ}\text{C}$
 $P_{tp} = 0.6117 \text{ kPa}$



At triple-point pressure and temperature, a substance exists in three phases in equilibrium.



(a) P - v diagram of a substance that contracts on freezing

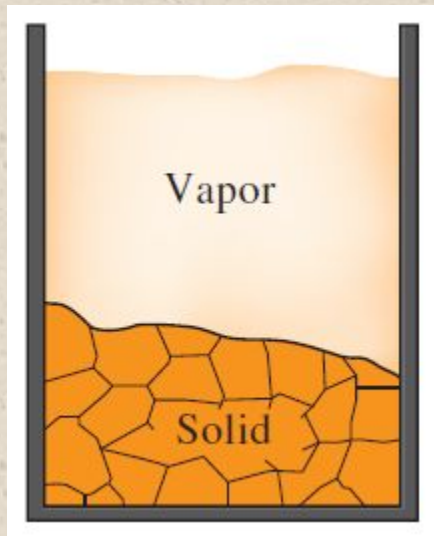


(b) P - v diagram of a substance that expands on freezing (such as water)

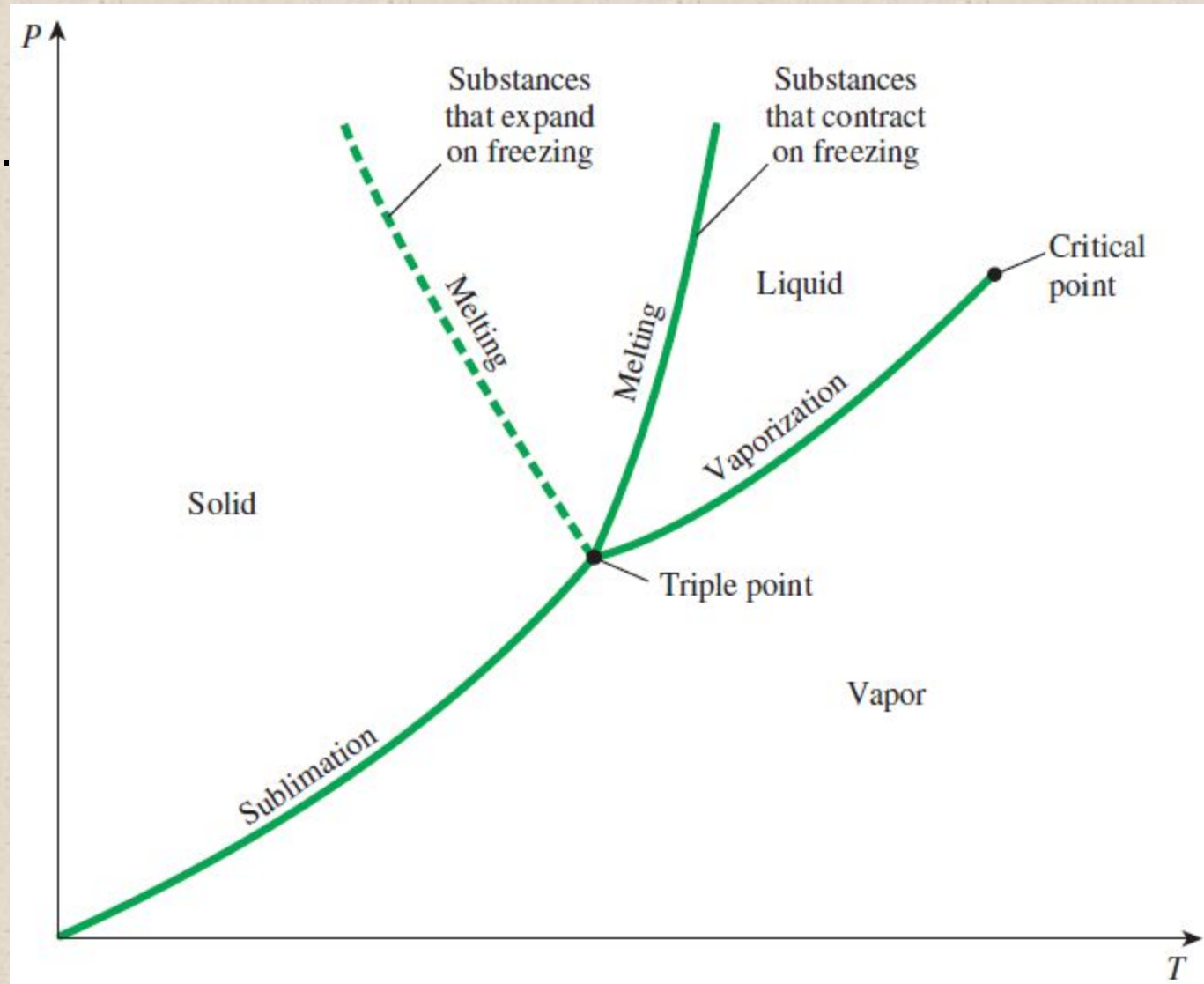
Phase Diagram

Sublimation:

Passing from the solid phase directly into the vapor phase.



At low pressures (below the triple-point value), solids evaporate without melting first (*sublimation*).



P-T diagram of pure substances.

The P - v - T surfaces present a great deal of information at once, but in a thermodynamic analysis it is more convenient to work with two-dimensional diagrams, such as the P - v and T - v diagrams.

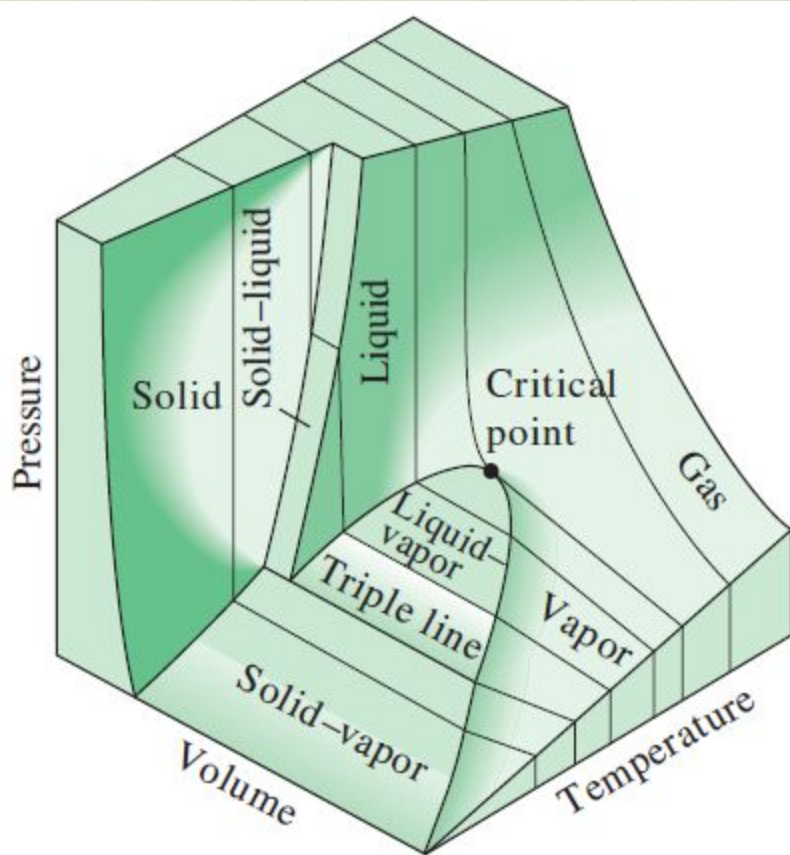


FIGURE 3-23

P - v - T surface of a substance that *contracts* on freezing.

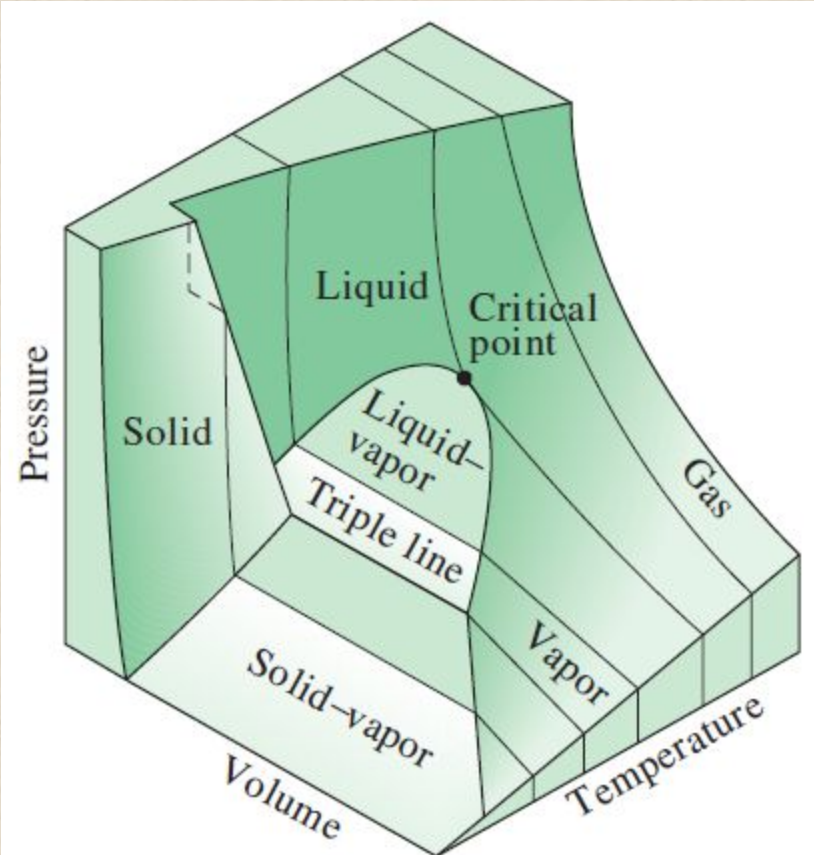


FIGURE 3-24

P - v - T surface of a substance that *expands* on freezing (like water).

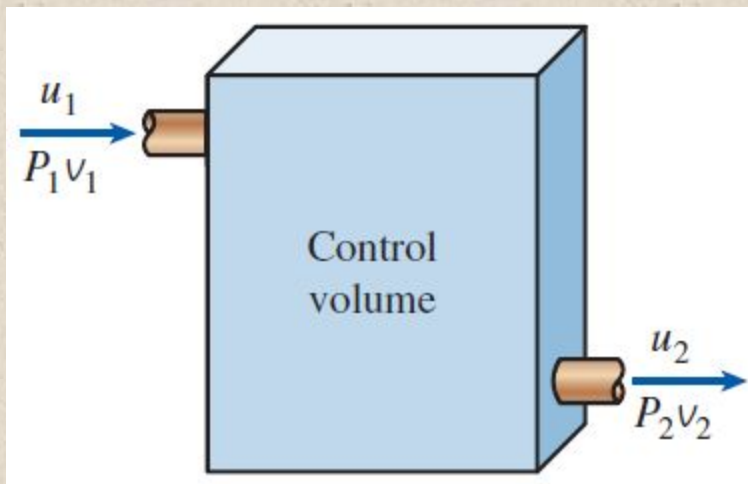
PROPERTY TABLES

- For most substances, the relationships among thermodynamic properties are too complex to be expressed by simple equations.
- Therefore, properties are frequently presented in the form of tables.
- Some thermodynamic properties can be measured easily, but others cannot and are calculated by using the relations between them and measurable properties.
- The results of these measurements and calculations are presented in tables in a convenient format.

Enthalpy—A Combination Property

$$h = u + Pv \quad (\text{kJ/kg})$$

$$H = U + PV \quad (\text{kJ})$$



The combination $u + Pv$ is frequently encountered in the analysis of control volumes.

$\text{kPa} \cdot \text{m}^3 \equiv \text{kJ}$
$\text{kPa} \cdot \text{m}^3 / \text{kg} \equiv \text{kJ/kg}$
$\text{bar} \cdot \text{m}^3 \equiv 100 \text{ kJ}$
$\text{MPa} \cdot \text{m}^3 \equiv 1000 \text{ kJ}$
$\text{psi} \cdot \text{ft}^3 \equiv 0.18505 \text{ Btu}$

The product *pressure* \times *volume* has energy units.

Saturated Liquid and Saturated Vapor States

A partial list of Table A-4.

v_f = specific volume of saturated liquid

v_g = specific volume of saturated vapor

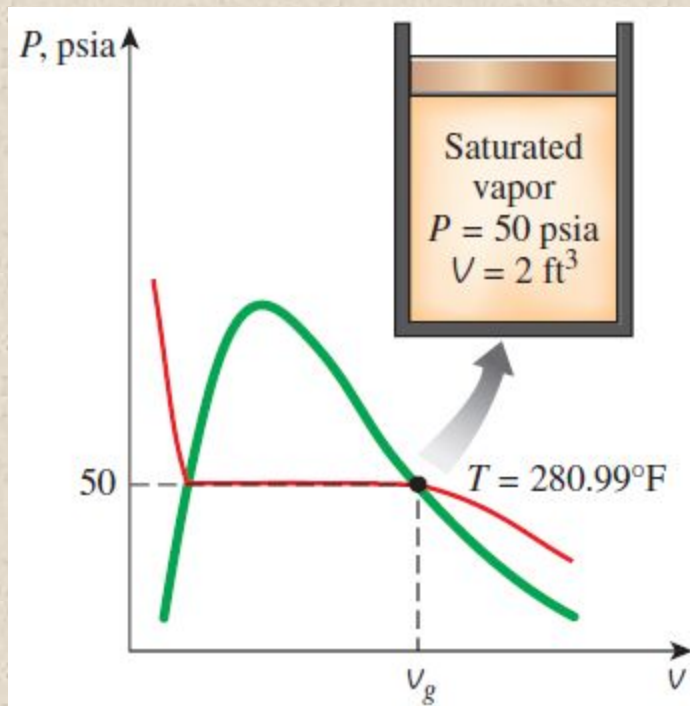
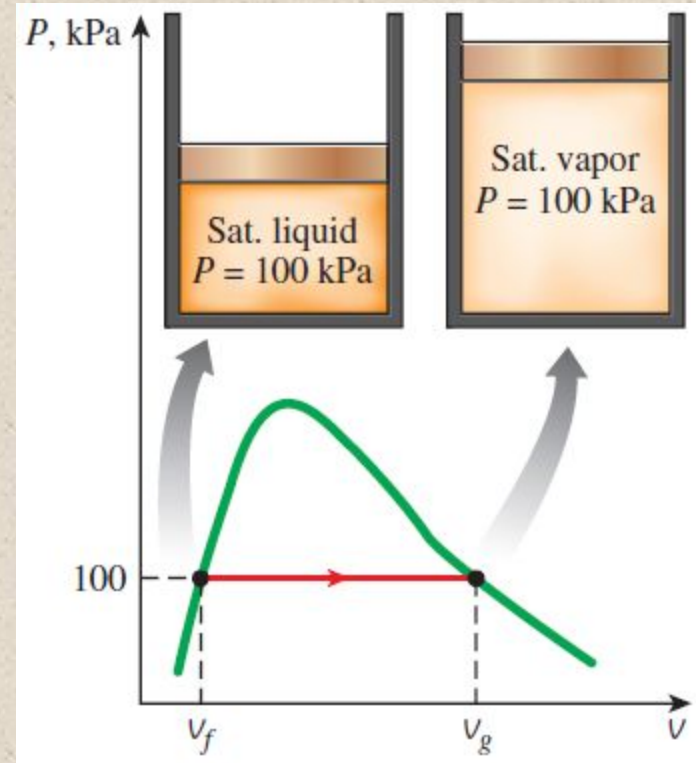
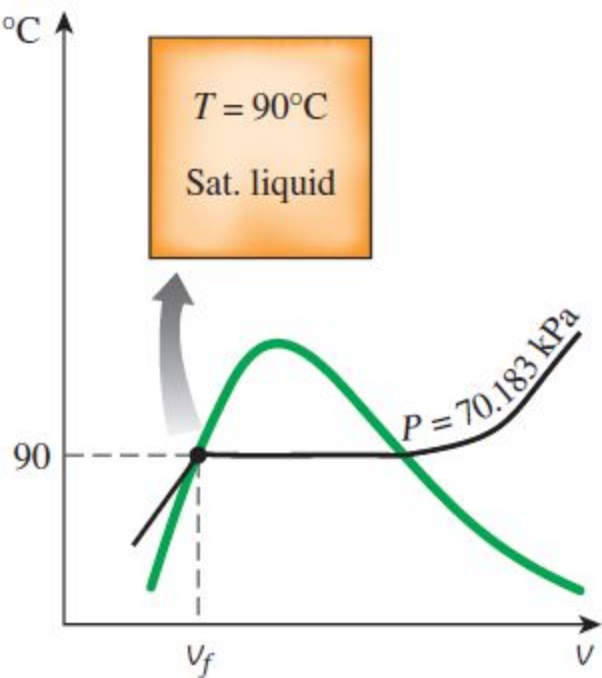
v_{fg} = difference between v_g and v_f (that is $v_{fg} = v_g - v_f$)

Temp. °C T	Sat. press. kPa P_{sat}	Specific volume m ³ /kg	
		Sat. liquid v_f	Sat. vapor v_g
85	57.868	0.001032	2.8261
90	70.183	0.001036	2.3593
95	84.609	0.001040	1.9808

- **Table A-4:** Saturation properties of water under temperature.
- **Table A-5:** Saturation properties of water under pressure.

Enthalpy of vaporization, h_{fg} (Latent heat of vaporization): The amount of energy needed to vaporize a unit mass of saturated liquid at a given temperature or pressure.

Examples:
Saturated liquid and saturated vapor states of water on T - v and P - v diagrams.



Saturated Liquid–Vapor Mixture

Quality, x : The ratio of the mass of vapor to the total mass of the mixture.

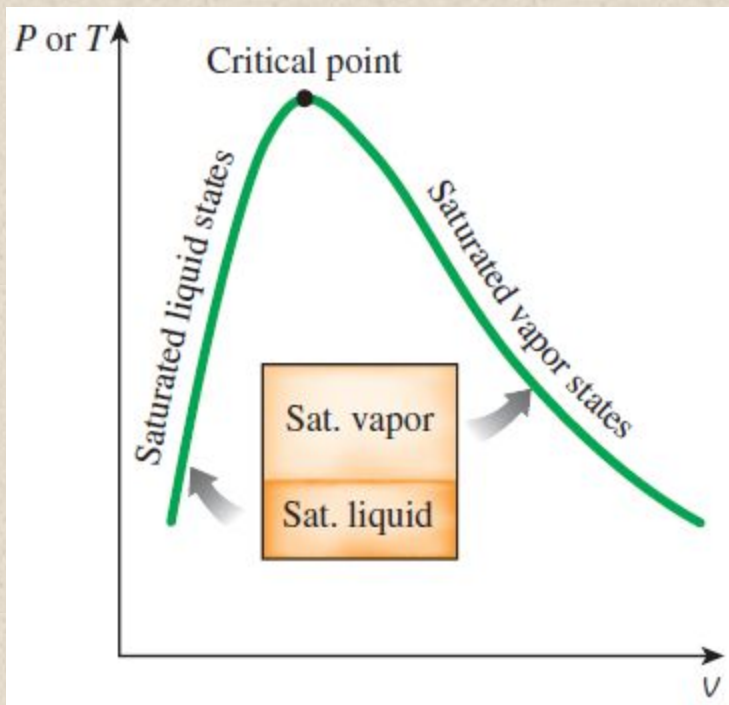
Quality is between 0 and 1 ➡ 0: sat. liquid, 1: sat. vapor.

The properties of the saturated liquid are the same whether it exists alone or in a mixture with saturated vapor.

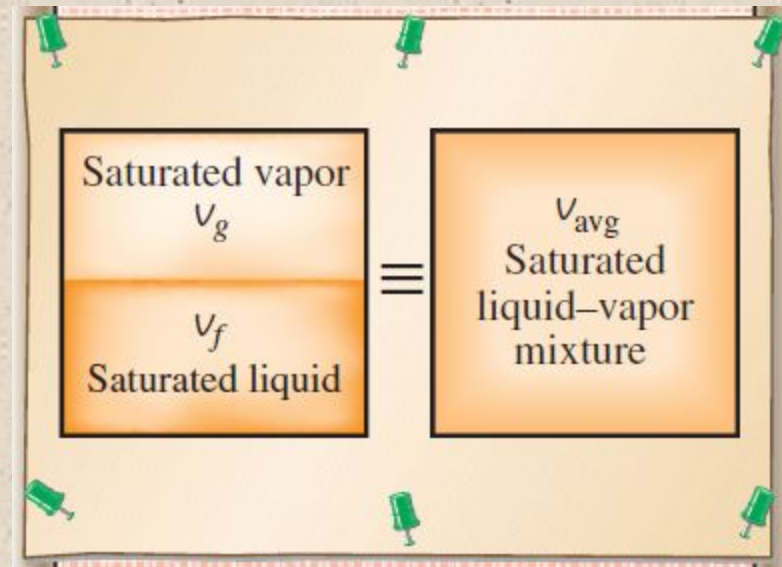
$$x = \frac{m_{\text{vapor}}}{m_{\text{total}}}$$

$$m_{\text{total}} = m_{\text{liquid}} + m_{\text{vapor}} = m_f + m_g$$

Temperature and pressure are dependent properties for a mixture.



The relative amounts of liquid and vapor phases in a saturated mixture are specified by the **quality x** .



A two-phase system can be treated as a homogeneous mixture for convenience.

$$v_{\text{avg}} = v_f + x v_{fg} \quad (\text{m}^3/\text{kg})$$

$$x = m_g/m_t \quad x = \frac{v_{\text{avg}} - v_f}{v_{fg}}$$

$$u_{\text{avg}} = u_f + x u_{fg} \quad (\text{kJ/kg})$$

$$h_{\text{avg}} = h_f + x h_{fg} \quad (\text{kJ/kg})$$

y ➡ v , u , or h .

$$y_{\text{avg}} = y_f + x y_{fg}$$

$$y_f \leq y_{\text{avg}} \leq y_g$$

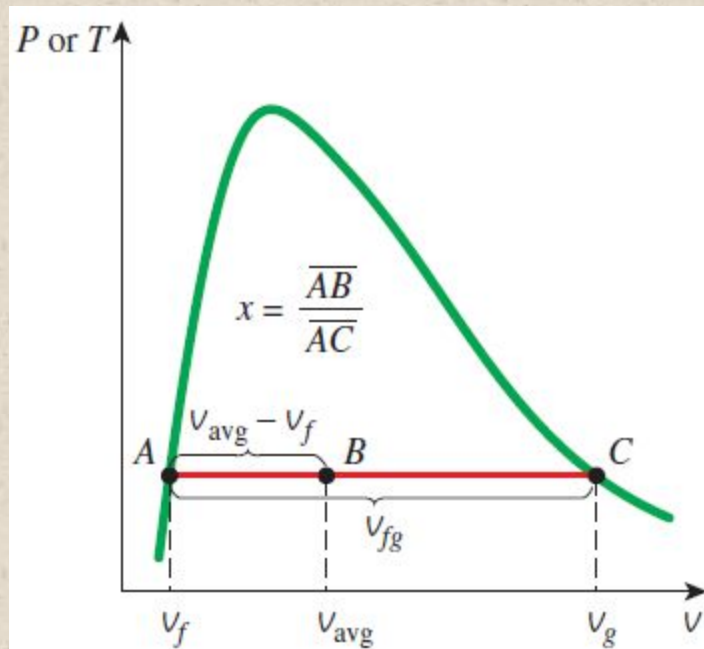


FIGURE 3–33

Quality is related to the horizontal distances on P - v and T - v diagrams.

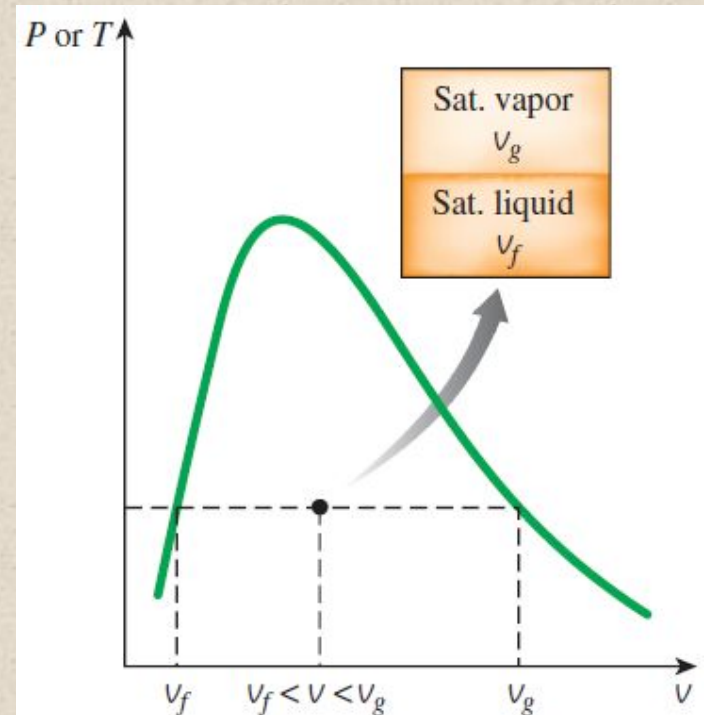


FIGURE 3–34

The v value of a saturated liquid–vapor mixture lies between the v_f and v_g values at the specified T or P .

Examples: Saturated liquid-vapor mixture states on T - v and P - v diagrams.

