

AMME2262 – SEMESTER 2

Engineering thermodynamics is the science of energy that applies to the design and analysis of energy conversion systems

Energy can be:

- Thermal or internal energy (stored in molecular forces)
- Chemical energy in molecular bonds (combustion)
- Kinetic energy (motion)
- Potential energy (elevation)
- Nuclear, magnetic and electric energy
- Sum of all energies = total energy E of the system

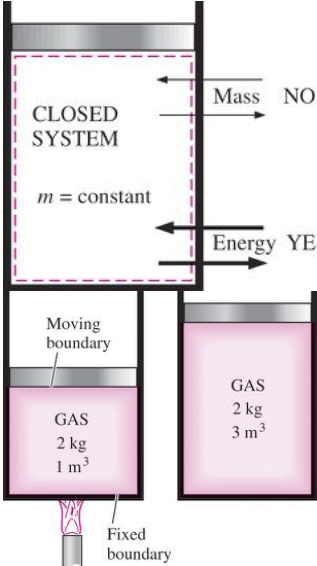
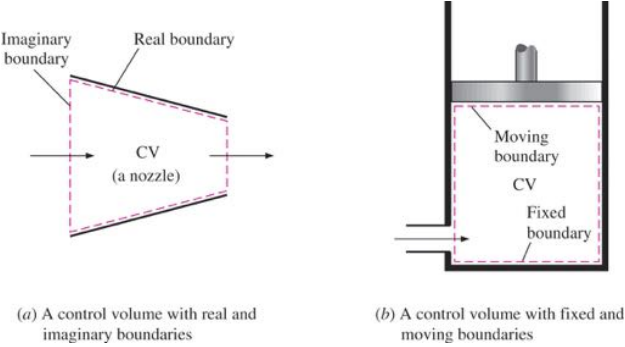
Important Terms:

Pressure (Pa, Nm⁻²)	<ul style="list-style-type: none"> • <u>Absolute pressure</u>: measured relative to absolute zero pressure (used mostly) • <u>Gage/Vacuum pressure</u>: measured relative to atmospheric pressure $P_{abs} = P_{atm} \pm P_{gage} \qquad P_{vac} = P_{atm} - P_{abs} \qquad P_{gage} = P_{abs} - P_{atm}$
Temperature (K)	<ul style="list-style-type: none"> • Use Kelvin Scale for units $T(K) = T(^{\circ}C) + 273.15$ <ul style="list-style-type: none"> • Engineering thermodynamics deals with temperatures in the range of 200K-3000K
Force (N)	<ul style="list-style-type: none"> • F=ma, or where a=g (gravity) F=mg • kgf, relative to local gravity
Pascal's law:	<ul style="list-style-type: none"> • The pressure applied to a confined fluid increases the pressure throughout by the same amount $P_1 = P_2 \rightarrow \frac{F_1}{A_1} = \frac{F_2}{A_2} \rightarrow \frac{F_2}{F_1} = \frac{A_2}{A_1}$

Density and specific gravity

Density (kgm⁻³)	Specific Volume (m³kg⁻¹)
$\rho = \frac{m}{V} \left(kg / m^3 \right)$	$v = \frac{V}{m} = \frac{1}{\rho} \left(m^3 / kg \right)$
Specific Weight (Nm⁻³)	Specific Gravity
$\gamma_s = \rho g$	$SG = \frac{\rho}{\rho_{H_2O}}$

Determining Type of System

Closed System	Open System
<ul style="list-style-type: none"> no mass transfer across boundaries Energy transfer across boundaries 	<ul style="list-style-type: none"> mass transfer across boundaries  <p>(a) A control volume with real and imaginary boundaries</p> <p>(b) A control volume with fixed and moving boundaries</p>

Assumption: Substances are treated as a **continuum** because of the very large number of molecules even in an extremely small volume (no need to consider on atomic level)

Properties of Systems:

Intensive Property	<p><i>Independent</i> of System Size</p> <ul style="list-style-type: none"> Temperature-T (K), Pressure-P (Pa), Density- ρ (kg/m³) 	<div style="border: 1px solid black; padding: 5px; text-align: center;"> m V T P ρ </div> <p style="text-align: center;">↓</p> <div style="display: flex; justify-content: space-around;"> <div style="border: 1px solid black; padding: 5px; text-align: center;"> $\frac{1}{2} m$ $\frac{1}{2} V$ T P ρ </div> <div style="border: 1px solid black; padding: 5px; text-align: center;"> $\frac{1}{2} m$ $\frac{1}{2} V$ T P ρ </div> </div> <div style="display: flex; justify-content: space-around; margin-top: 5px;"> } Extensive properties } Intensive properties </div>
Extensive Property	<p><i>Dependent</i> of System Size</p> <ul style="list-style-type: none"> Mass - m (kg), Volume - V (m³), Energy - E (J) 	
Specific Properties	<p>Extensive Properties per unit mass that become intensive properties</p> <ul style="list-style-type: none"> mass-m (kg) $V/m=$, specific volume (m³/kg) $U/m=u$, specific internal energy (kJ/kg) $E/m=e$, specific energy (kJ/kg) 	

Units (metric system)

All equations must be dimensionally homogeneous

Equilibrium

- System in thermodynamic equilibrium if it is in thermal, mechanical, phase and chemical equilibrium.
- Properties of system do not change over entire system

Temperature and the Zeroth law of thermodynamics

- Zeroth law states that two bodies are in thermal equilibrium if both have the same temperature reading even without being in contact
- Heat moves from hot to cold objects

Equations Of State (The Ideal Gas Equation)

Ideal Gas Equation relates pressure, temperature and specific volume of a substance. The assumptions made for it to hold true include:

- The gas is composed of a large amount of small molecules
- The gas molecules are elastic (KE is conserved)
- The size and total volume of the molecules is small relative to the volume
- Thermal motions of the gas are random

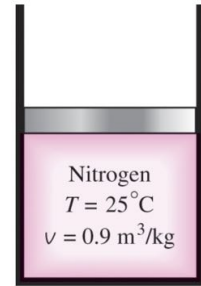
Note: real gases can be approximated by an ideal gas at “low” densities: “low” pressures and “high” temperatures

Substance	R, kJ/kg.K
Air	0.2870
Helium	2.0769
Argon	0.2081
Nitrogen	0.2968

State Postulate

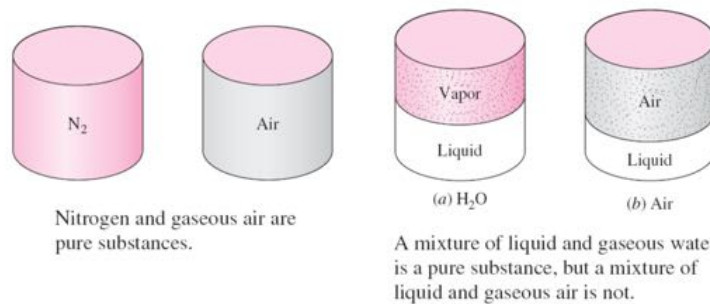
State postulate: The number of properties required to fix the state of a system

- The state of a **simple* compressible system** (i.e. A system that involves no electrical, magnetic, gravitational, motion, and surface tension effects) is completely specified by **2 independent, intensive properties**
- The state of the nitrogen here is fixed by two independent, intensive properties

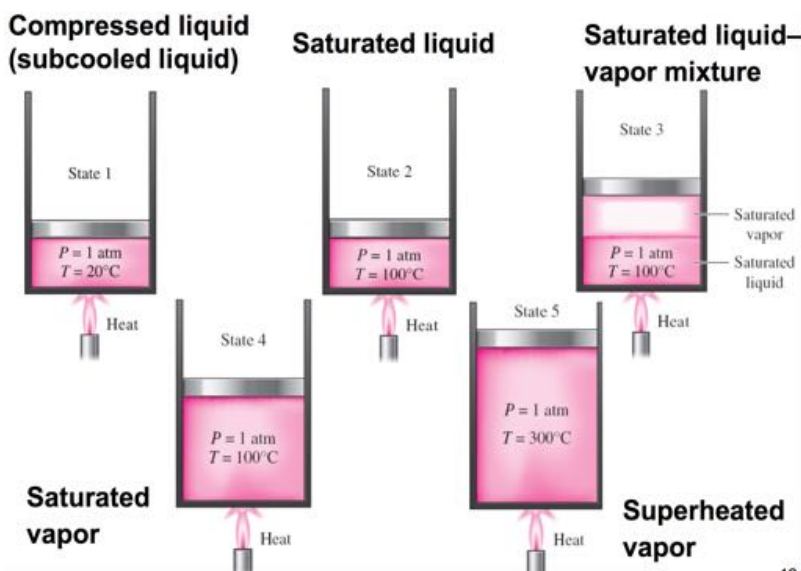


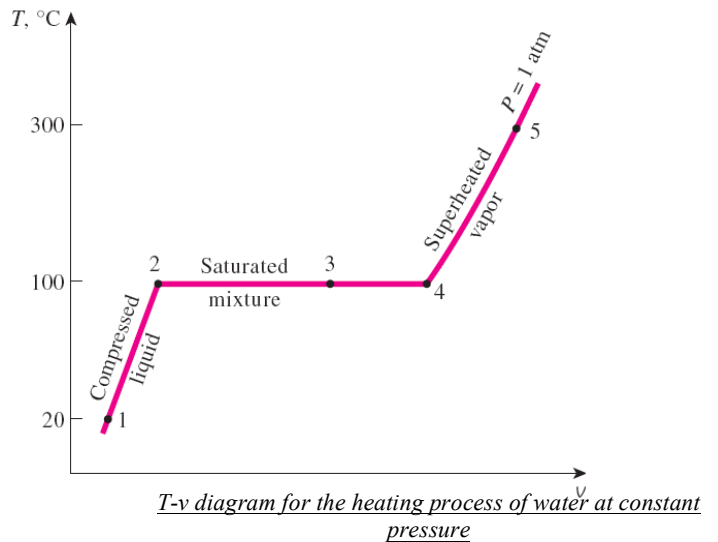
Pure Substances

- A substance that has a fixed chemical composition throughout.
- Air is a pure substance at atmospheric pressure and temperature, despite being a mixture of several gases



Phase Change	Description
1. Compressed liquid (subcooled liquid):	A substance that it is not about to vaporize (i.e. liquid phase)
2. Saturated liquid:	Exists as liquid that is about to vaporize.
3. Saturated liquid–vapor mixture:	The state at which the <i>liquid and vapor phases coexist</i> in equilibrium
4. Saturated vapor:	A vapor that is about to condense
5. Superheated Vapour	A vapor that is not about to condense (i.e., not a saturated vapor)

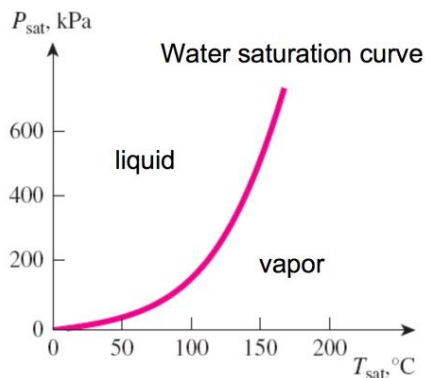




• **Reversing** the process 5 → 1 following the same path releases the **same amount of heat** as was added in the forward process. The temperature at which water starts boiling depends on the pressure; therefore, **if the pressure is fixed, so is the boiling temperature.**

Saturation temperature T_{sat} : The temperature at which a pure substance changes phase at a given pressure.

Saturation pressure P_{sat} : The pressure at which a pure substance changes phase at a given temperature.



The liquid–vapor saturation curve of a pure substance (water) – water boils at 100°C at 1 atm pressure

Saturation (boiling) pressure of water at various temperatures

Temperature, $T, ^\circ\text{C}$	Saturation pressure, P_{sat}, kPa
-10	0.26
-5	0.40
0	0.61
5	0.87
10	1.23
15	1.71
20	2.34
25	3.17
30	4.25
40	7.39
50	12.35
100	101.4
150	476.2
200	1555
250	3976
300	8588

Variation of the standard atmospheric pressure and the boiling (saturation) temperature of water with altitude

Elevation m	Atmospheric pressure, kPa	Boiling temperature, °C
0	101.33	100.0
1,000	89.55	96.5
2,000	79.50	93.3
5,000	54.05	83.3
10,000	26.50	66.3
20,000	5.53	34.7

<p>At a given temperature if:</p> <ul style="list-style-type: none"> • Pressure < liquid-vapor saturation pressure = vapor • Pressure > liquid-vapor saturation pressure = liquid 	<p>At a given pressure if:</p> <ul style="list-style-type: none"> • Temp. < liquid-vapor saturation temp. = liquid • Temp. > liquid-vapor saturation temp. = vapour
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Latent Heat

Latent heat:	amount of energy absorbed or released during a phase- change process
Latent heat of fusion:	amount of energy absorbed during melting (equal to energy released during freezing)
Latent heat of	amount of energy absorbed during vaporization (equal to energy released

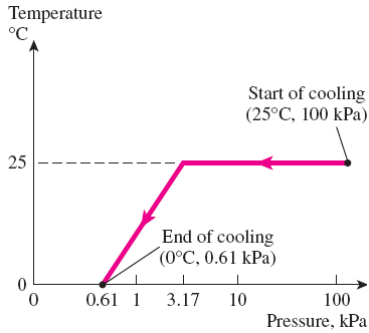
vaporization:

during condensation)

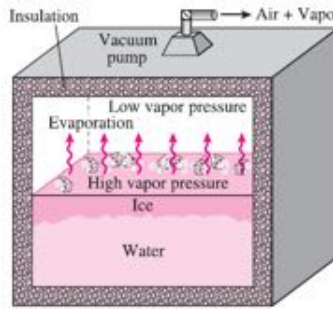
*Magnitudes of the latent heats dependent on temp. or pressure at which the phase change occurs.

e.g. At 1 atm pressure, the latent heat of fusion of water = 333.7 kJ/kg and the latent heat of vaporization = 2256.5 kJ/kg.

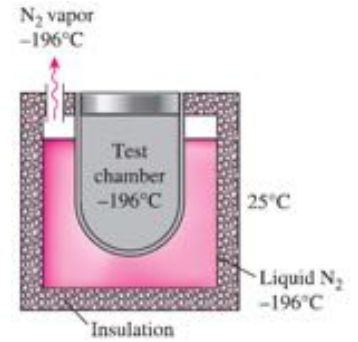
Consequences of T_{sat} and P_{sat} Dependence



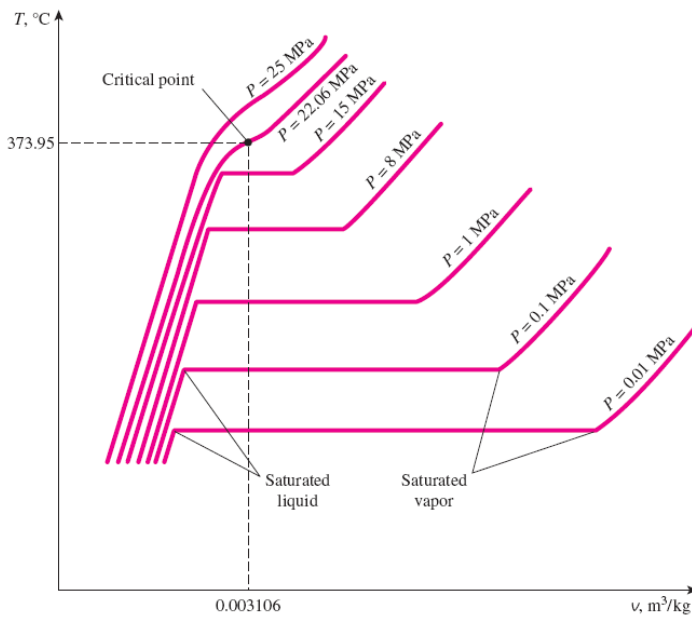
The variation of the temperature of fruits and vegetables with pressure during vacuum cooling from 25°C to 0°C



Ice made by evacuating air space in water tank



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



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