# ENERGY BALANCES



#### Energy was first hypothesized by Newton.

# Newton was the first who expressed kinetic and potential energies.



Potantial,
 Kinetic,
 Chemical,
 Mechanical,

Electrical

#### Energy cannot be measured directly.



# Energy of an object due to its <u>height</u>. P.E. = m g h







 Atoms and molecules move constantly in random direction. Atoms vibrate, rotate and collide with each other.

Energies related to all these movements are called as the internal energy.

# During processing of food;

One or two forms of energy are predominant, the others can be omitted.

## Example-1

Depositing of sugar beets from conveyor to the tank



- Potantial and kinetic energy changes,
- Chemical end electrical energies do not change





# During heating of tomato juice in a heat exchanger,

- Potantial and kinetic energy do not change,
  - Internal energy changes as a result of change in temperature





Not static, always in flux.

- Even under steady-state conditions, an object absorbs energy from its surroundings and at the same time emits energy to its surroundings at the same rate.
- When there is imbalance between the energy absorbed and emitted, the steady-state is altered, molecular energy in the system may increase or new compounds may be formed.

# First law of thermodynamics (law of conservation of energy);

In an isolated system, <u>overall energy</u> <u>does not change.</u>

Indicating that the energy <u>do not</u> form or <u>disappear</u>.

Only the energy in one form can be <u>converted</u> to another form.

# **Energy balance equation:**

#### determined as in the case of mass balances.





#### Accumulation will be zero.



# If the system involves only an exchange of energy between two components;

Energy gain	Energy loss
by	by
component	component
1	2

# Energy balances are done for;

Identifying areas where energy conservation can be done.

In designing heating or cooling systems to ensure the right amount of fluids used for heat exchange and the right size of equipment used to achieve the processing objectives.



#### Sum of **internal energy** and the **multiplication** of pressure and specific volume H = E + PV'

- H: Enthalpy, kJ/kg
- E: Internal energy, kJ/kg
- P: Pressure, kPa
- V': Specific volume, m<sup>3</sup>/kg

When the enthalpy of an air in a room is described, then this enthalpy includes the internal energy.

If a liquid enters and leaves the open system, then the enthalpy will be equal to the multiplication of pressure and specific volume (flow energy).



- Enthalpy cannot be measured directly.
- Enthalpy values should always be given on the reference conditions (in particular temperature).
- For steam tables, reference temperature is 0°C.



A form of energy.

 "Heat exchange occurs between a system and its surroundings because of <u>temperature difference</u>.

Heat transfers from a hot object to a cold one.

Heat is used in many food applications, particularly in cooking and preservation (pasteurization and sterilization).

Common heat units are calorie, kilo-calorie, BTU and joule.

# Heat (Q);

#### The sign of Q shows the flux of heat.

If heat transfers from a system to its surroundings



if heat transfers from the surroundings into a system (heating of potato)



# **Specific heat**

- Amount of heat gained or lost for a unit mass of a substance, accompanying <u>a unit</u> <u>change in temperature</u> of the substance but <u>not involving with phase change.</u>
- Specific heat of solids and liquids are constant over a fairly wide temperature range.
- Specific heat of gases varies with temperature.



•  $c_p$  and  $c_v$  are similar for solids and liquids.

•  $c_p$  and  $c_v$  are different for gases.

Since most food processes are carried out under constant pressure, the specific heat values under constant pressure are used for food processing calculations.

### Forms of heat



#### Sensible heat;

energy associated to change in the temperature of system or object (food)

#### Laten heat;

energy associated with phase transitions (no temperature changes)

## **Sensible heat of foods**

 $Q = m C_p (\Delta T)$ 

- Q : Sensible heat gained by food, kJ
- m: Mass of food, kg
- C<sub>p</sub>: Specific heat of food, kJ/kg °C
- $\Delta$  T : Change in temp of food, °C

Example 4.1: Calculate the enthalpy change during heating of 3 kg water from 20°C to 90°C.

#### **Sensible and latent heat:**



If we continue to heat water, temperature of water will also increase and finally water boils (Sensible heat).

If we still heat <u>boiling</u> water, water molecules will start to transfer to vapor phase. During this process, temperature of vapor will still be 100°C (just phase transition). In order to evaporate the water molecules, we need to give the <u>latent heat of vaporization</u>.

Latent heat of condensation is equal to latent heat of evaporation.

Condensation is the phase change from vapor phase to liquid phase.

 If heat is removed from vapor, then distance between vapor molecules will decrease and vapor molecules will condensate in droplets. Heat removed during this process is called as latent heat of condensation.

- If water in liquid state changes to solid phase, heat released will be *latent heat of freezing*.
- During <u>melting</u> of a frozen material, the same amount of heat absorbed and this heat is called as *latent heat of fusion*.

- Boiling point of water depends on the pressure.
- At 1 atm, water boils at 100°C.
- If pressure is increased, then boiling point will also increase.
- Latent heat of vaporization of water also depends on boiling point of water.
- The relationship between pressure, boiling water and <u>latent heat of vaporization</u> is given in Table 4.1.

Pressure (atm)	Boling point of water (°C)	<i>Latent heat of vaporization</i> (kcal/kg)
1	100	540
1.5	110	532.5
0.2	60	563

Example 4.2: 3 kg of water at 20°C was boiled under 1.5 atm of pressure and 110°C, and then transformed to vapor. Calculate the enthalpy change for the vapor 110°C. Example 4.3: The temperature of 2 kg water at 20°C increases to 110°C after heating under 1.5 atmosphere constant pressure. After continuous heating, the temperature of the resulting vapor at 110°C increases to 125°C. The specific heat of vapor is <u>0.445 kcal/kg °C</u>.

- a) Calculate the enthalpy change during transforming 2 kg water to vapor at 125°C.
- b) Calculate the entalphy of vapor at 125°C.

- Final enthalpy (629.2) is not the enthalpy of vapor at 125°C.
- It is the total heat needed to transfer 1 kg water to 1 kg vapor at 125°C.
- If we want to calculate enthalpy of vapor at 125°C, we need to take into consideration of enthalpy of water between 0° and 20°C. The reference point in enthalpy calculation is 0°C. Q = (1 kg) (1 kcal/kg °C) (20 0)°C = 20 kcal
  Enthalpy of vapor at 125°C will be the sum of "20 + 629.2 = 649.2 kcal."

Example 4.4: The ice at -10°C was melted by heating and then under normal atmospheric pressure, the resulting water molecules were transferred to vapor at 100°C. Calculate the enthalpy change during transforming 5 kg ice to vapor. The specific heat of ice is 0.5 kcal/kg °C and *latent heat of fusion* is 80 kcal/kg.

**Example 4.5:** The fruit at 25°C is frozen to – 20°C. The specific heat of fruit above freezing point is 0.95 kcal/kg °C and the freezing point of fruit is -3°C, the specific heat of fruit below freezing point is 0.460 kcal/kg °C and latent heat of freezing is 68 kcal/kg. Calculate the enthalpy change during freezing of 8 kg fruit from  $25^{\circ}$ C to  $-20^{\circ}$ C.

![](_page_38_Picture_0.jpeg)

Example 4.6: 6 kg of water at 25°C was boiled to 104°C under 1.1 atm of pressure. And then, the steam was heated to 130°C.

- a) Find out the enthalpy change in "kcals" during the heating of water at 25°C to vapor at 130°C.
- b) Find out the enthalpy of vapor at 130°C in SI unit system by taking into consideration of enthalpy calculations for water.

## Data

- latent heat of evaporation of water; 540 kcal/kg at 100°C, 532.5 kcal/kg at 110°C.
- specific heat of vapor, 1886 J/kg K at 125°C and 1891 J/kg K at 150°C.

![](_page_41_Picture_0.jpeg)

#### E = 3766 kcal (entalphy change from 25°C to 130°C)

#### E<sub>130°C</sub>= 3916 kcal (entalphy at 130°C)

![](_page_42_Picture_0.jpeg)

Amount of heat gained or lost for a unit mass of a substance, accompanying <u>a unit change in</u> <u>temperature</u> but <u>not involving with phase</u> <u>change.</u>

![](_page_42_Picture_2.jpeg)

Most solids and liquids have a constant specific heat over a fairly wide temperature range above freezing point.

Specific heat of gases varies with temperature.

# **Specific heat of foods;**

Depends on <u>composition</u> and <u>mostly water</u> <u>content</u>.

- Since the food composition changes depending on many factors, most accurate way of calculating specific heat of foods is to do <u>experiment</u>.
- In practice, specific heat of foods is calculated using the equations, if the composition of foods (mass fractions) is known,

- One of the most important compositional compound of foods is <u>water</u>.
- Specific of heat of water in liquid and solid phase is very <u>different.</u>

#### Therefore, specific heat of foods,

Above freezing point

Below freezing point, are very <u>different.</u>

# Above freezing point (Siebel equation)

$$C_p = 1 m_s + 0.2 (1 - m_s)$$

- m<sub>s</sub>: Mass fraction of water in food
- 1: Specific heat of water, kcal kg<sup>-1</sup>°C<sup>-1</sup> or BTU  $lb_m^{-1}°F^{-1}$
- **0.2**: Specific heat of nonfat solids, kcal kg<sup>-1</sup>°C<sup>-1</sup> or BTU  $Ib_m^{-1}°F^{-1}$
- 1 m<sub>s</sub> : Mass fraction of nonfat solids in food containing no fat
- C<sub>p</sub>: Specific heat, kcal kg<sup>-1</sup> °C<sup>-1</sup> or BTU lb<sub>m</sub><sup>-1</sup> °F<sup>-1</sup>

# **Arranging the equation:**

$$C_{p} = 1 m_{s} + 0.2 - 0.2 m_{s}$$
$$C_{p} = m_{s} - 0.2 m_{s} + 0.2$$
$$C_{p} = (1 - 0.2) m_{s} + 0.2$$

 $C_p = 0.8 m_s + 0.2$  (English, mks and cgs)

![](_page_48_Figure_0.jpeg)

- C<sub>p</sub> : Specific heat above freezing point, kJ kg<sup>-1</sup> K<sup>-1</sup>
- m<sub>s</sub> : Mass fraction of water in food

# Effect of fat content on specific heat:

For the animal based products which contain considerable <u>amount of fat</u>, the specific heat of food should be estimated by taking into consideration of the <u>mass</u> <u>fraction of fat (F)</u>. Specific heat of foods containing fats above freezing point

 $C_p = 0.4 F + 0.2 NFS + 1 M$  (English, mks and cgs)

 $C_p = 1.67472 F + 0.83736 NFS + 4.1868 M$  (SI)

- **C**<sub>p</sub>: Specific heat of food containing considerable amount of fat above freezing point, kJ/kg K
- F: Mass fraction of fat in food
- NFS : Mass fraction of non-fat solid in food
  - M : Mass fraction of water in food

# Taking into consideration of carbohydrate, protein and ash:

# $C_{p \text{ (food)}} = 1.424 \text{ C} + 1.549 \text{ P} + 1.675 \text{ F} + 0.837 \text{ A} + 4.187 \text{ M}$

- C<sub>p (food)</sub> : <u>kJ/kg K</u>
- Note: (1) C, P, F, A and M is denoted for carbohydrate, protein, fat, ash and water, respectively.

(2) Numbers in front of mass fractions of food constituents is the actual specific heat of each constituent.

Example 4.7: Calculate the specific heat of orange juice concentrate having a solid content of 45% in SI unit system.

## Answer

Example 4.9: Calculate the specific heat of beef roast containing 15% protein, 20% fat and 65% water in SI unit system.

## Answer

• 
$$C_{\text{roast beef}} = 0.76 \text{ kcal/(kg °C)}$$
  
•  $C_{\text{roast beef}} = 3181.4 \text{ J/(kg K)}$ 

Example 4.10: Calculate the specific heat of food containing 30% of carbohydrate, 10% of protein, no fat, 0.4% of ash, 56% of water in SI unit system.

#### Answer

# $C_{food} = 1.424 C + 1.549 P + 1.675 F + 0.837A + 4.187 W$

#### $C_{food} = 1.424 (0.3) + 1.549 (0.1) + 0.837 (0.004)$ + 4.187 (0.56)

• 
$$C_{food} = 2.93 \text{ kJ/(kg K)}$$