#### **Basics of bioenergetics**

**Medical Chemistry** 

#### Definitions

Thermodynamics studies transformations of energy.

A thermodynamic system is an object or a group of interacting objects separated with a boundary from the surroundings



**Homogeneous and** heterogeneous thermodynamic systems Homogeneous systems consist of a single phase They do not have a phase interface.

Example: water solutions of glucose, sucrose, sodium chloride etc.



**Homogeneous and** heterogeneous thermodynamic systems Heterogeneous systems consist of several phases They have a phase interface. Examples:









Open systems exchange both matter and energy with the surroundings

Examples:



Closed systems exchange energy and do not exchange matter.

#### **Examples:**



Isolated systems exchange neither mass nor energy with the surrounudings.

Example: a thermocapsulated cup



**Thermodynamic potentials** Internal energy, U Enthalpy, H Entropy, S Gibbs energy, G

Internal energy, U Internal energy of a system is the sum of the potential energy and the kinetic energy of all components of the system. In it, the kinetic is the energy of molecular movement. The potential energy is the energy in all the chemical bonds. OH H - C - H



#### Internal energy, U

Internal energy excludes kinetic energy due to motion of the system as a whole and gravitational potential energy due to its elevation.



Internal energy cannot be measured but its change ΔU can be measured

#### Enthalpy, H

Enthalpy is the total heat content of the system.

Enthalpy is the sum of internal energy and the work done by the system to stay in its boundaries:

#### H = U + pV

Enthalpy cannot be measured, but its change ΔH can be measured.

# Enthalpy, HΔH is the heat effect of a reaction.ΔH < 0: heat is released,</td>the reaction is exothermic



CARBON DIOXIDE

**ATER** 

SUGARS

WATER

**ΔH > 0: heat is absorbed, the reaction is endothermic** 

# Standard enthalpy of formation

Standard enthalpy of formation of a compound is the enthalpy change that occurs when one mole of a substance in its standard state is formed from the component elements in their standard states.



Standard enthalpy of formation of elements (simple substances) is equal to zero at standard conditions. Examples of simple substances: O<sub>2</sub>, N<sub>2</sub>, Mn, Fe, Br<sub>2</sub>, K, C, etc.

#### Standard enthalpy of combustion

Standard enthalpy of combustion is of a compound is the enthalpy change that occurs when 1 mole of a substance reacts completely with oxygen under standard conditions.



#### Hess' Law

The heat effect of a reaction depends only on the initial and final states of the system, and is independent of the pathway between these states.



#### $\Delta H_1 = \Delta H_2 + \Delta H_3 = \Delta H_4 + \Delta H_5 + \Delta H_6$

#### Hess' Law

Hess law is a definition of the First Law of Thermodynamics.

It allows ΔH for a reaction to be calculated even when it cannot be directly measured.



Application: find energy values of food

**First Law of Thermodynamics** Definition 1: Law of Conservation of Energy

Energy can neither be created nor destroyed, it can only be transformed from one form to another First Law of Thermodynamics Definition 2: In an isolated system, the sum of all types of energy is a constant value First Law of Thermodynamics

Definition 3: Heat transferred to a system is used to do work and change internal energy of the system.

#### $Q = \Delta U + A$

Q is the heat exchanged between a system and its surroundings;

AU is the total change in internal energy of a system;

A is the work done by or on the system

# First Law of Thermodynamics

#### **Definition 4:**

The perpetuum motion machines of the first kind are impossible.

#### No Energy = No Work



#### **Definition 1:**

## Entropy is the measure of randomness (disorder).



#### Entropy, S

#### **Definition 2:**

## Entropy is the energy that cannot be converted into work.





#### Entropy, S

#### **Definition 3:**

#### Entropy is the dissipated (wasted) energy



## **Entropy, S** Entropy increases from solid to liquid to gas.



Entropy increases with an increase in temperature.



Entropy of a system and its surroundings increases in spontaneous processes

Entropy increases with time.





## A change of entropy $\Delta S$ is the measure of irreversibility of a process





Second Law of Thermodynamics Definition 1:

It is impossible to create a machine that can transform all heat into useful work.

Second Law of Thermodynamics Definition 2:

All energy tends to transform into heat, and the heat tends to dissipate Second Law of Thermodynamics Definition 3:

Heat cannot flow spontaneously from a material at a lower temperature to a material at a higher temperature.

Second Law of Thermodynamics Definition 4:

In an isolated system, entropy increases and reaches its maximum at equilibrium

i.e.

## No work is done at equilibrium and

Living organisms are non-equilibrium systems Second Law of Thermodynamics Definition 5: Entropy is the arrow of time.

#### Gibbs energy, G

- 1. Gibbs energy is free energy
- 2. Gibbs energy is the energy that can be used to do work

#### $\mathbf{G} = \mathbf{H} - \mathbf{T}\mathbf{S}$

H is enthalpy T is absolute temperature S is entropy

### **Gibbs energy change,** $\Delta G$ $\Delta G = \Delta H - T\Delta S$ $\Delta G$ is the only criterion of spontaneous

- processes
- **ΔG < 0: the process is spontaneous** (exergonic)
- **ΔG > 0:** the process is not spontaneous (endergonic)
- $\Delta G = 0$ : the process is in equilibrium

**Endergonic and exergonic** processes in the organism **Exergonic processes: glycolysis, fatty** acid oxidation, ATP hydrolysis **Endergonic processes: synthesis of** glucose, fatty acids, proteins, ATP Exergonic and endergonic processes are conjugated, i.e. an endergonic process only proceeds with energy from an exergonic process.