

Gibbs Energy

Gibbs energy (also known as *free enthalpy*, *Gibbs free energy*, *Gibbs function* or *Gibbs potential*)

is a state quantity, usually marked with the letter **G**, representing that part of the heat content of the system that can be used to convert it into another form of energy. The rest of the thermal content of the system can only be converted into heat. Its unit in the SI system is the joule (J).

The reason for its introduction was that entropy as a **criterion of the arbitrariness of the process** is only suitable for adiabatic processes. The newly introduced thermodynamic potential - Gibbs free energy, represents a criterion suitable for more general conditions (for isothermal - isobaric processes).

The free energy (in a system with a constant number of particles) can be expressed by the relation:

$$G = H - TS,$$

where **H** is the total **heat content of the system**, **T** is the thermodynamic temperature (K) and **S** is the state quantity **entropy**. It follows from the given relationship that the smaller the entropy value, the greater the amount of Gibbs free energy the system has.

Living organisms thus obtain the most usable energy from structurally complex compounds such as polysaccharides, proteins and lipids, which have a low entropy value. On the contrary, a large amount of energy is consumed during the synthesis of these compounds.

Enthalpy H consists of the Gibbs energy (free enthalpy) **G** and the bound energy that cannot be converted into work at a given temperature.

After substituting for the enthalpy **H** we get:

$$G = U + pV - TS$$

where **U** is the internal energy, **p** is the pressure, **V** is the volume within the system, **T** is the thermodynamic temperature and **S** is the entropy of the system.

During chemical transformations the Gibbs energy content changes between the initial state G1 and the final state G2 of the system:

Failed to parse (SVG with PNG fallback (MathML can be enabled via browser plugin): Invalid response ("Math extension cannot connect to Restbase.") from server "https://wikimedia.org/api/rest_v1/"): $\Delta G = G_2 - G_1$

Three situations can occur:

a) $G_2 < G_1$ and therefore **Failed to parse (syntax error): $\Delta G < 0$**

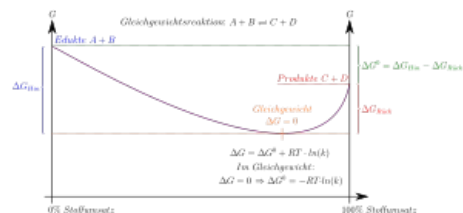
In this case, the Gibbs energy of the products is smaller than the Gibbs energy of the substrates, energy is released and it is therefore an exergonic process. This energy can be further converted into work or another type of energy, in the cell, for example, for ATP biosynthesis. In the organism, this situation occurs in the case of catabolic reactions, during which complex compounds are degraded and energy is released.

b) $G_2 > G_1$ and therefore **Failed to parse (syntax error): $\Delta G > 0$**

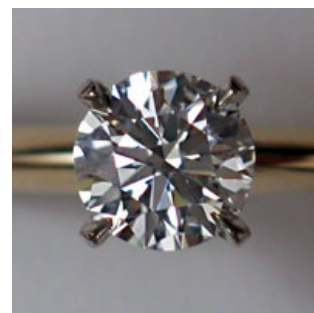
The Gibbs energy of the products has a higher value than the energy of the substrates, so it is an endergonic reaction, in which we have to supply the energy ΔG to the system. Anabolic processes take place in the body, during which complex structures are synthesized, or active transport of substances through the cell membrane.

c) **Failed to parse (syntax error): $\Delta G = 0$**

In this case, there is no change in the energy content of the system, the system is in equilibrium.



Change in Gibbs energy during an equilibrium reaction depending on the degree of progress of the chemical reaction. A complete transformation of the starting substances will not occur, because the reverse reaction of the products works against it. Chemical equilibrium occurs for such a concentration of products at which $\Delta G = 0$.



The reaction $C(s)_{\text{diamond}} \rightleftharpoons C(s)_{\text{graphite}}$ takes place at 25°C and 1 atm. to a negative change in the Gibbs free energy, and is therefore a thermodynamically spontaneous process. Graphite is a more stable form of carbon. The transformation of diamond into graphite takes place spontaneously, but it takes a very long time. The fact that the event is thermodynamically arbitrary does not tell how long it will take for the equilibrium to be established. Thermodynamics does not solve the problem of time.

Links

Literature

- MATOUŠ, Bohuslav. *Základy lékařské chemie a biochemie*. 1. edition. Galén, 2010. 540 pp. ISBN 978-80-7262-702-8.
- KOLEKTIV, Autorů. *Gibbs free energy* [online]. [cit. 2014-12-08]. <https://en.wikipedia.org/wiki/Gibbs_free_energy>.
- BOJKOVSKÝ, Martin. *Gibbsova energie* [online]. [cit. 2014-12-08]. <http://fikus.omska.cz/~bojkovsm/termodynamika/gibbsova_energie.html>.

Related Articles

- Free Energy

External Links

- Co je Gibbsova energie? článek na serveru Česká chemie.cz (http://www.ceskachemie.cz/svet-chemie/popularni-chemie/chemicka-poradna/co-je-gibbsova-energie#.Uvkfh_I5M3U)
- Gibbsova energie, RNDr. Martin Bojkovský, 2009 (http://fikus.omska.cz/~bojkovsm/termodynamika/gibbsova_energie.html)
- <https://de.wikipedia.org/wiki/Gibbs-Energie>