

## Thermodynamic modeling of biological growth

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### Abstract

A few different approaches have been utilized as a part of an endeavor to characterize and dissect the thermodynamics modeling of biological growth in bioreactor culture. While thermodynamic theory has been created adequately to empower attractive forecast of biomass and catabolic-item yield, expectation of non-catabolic item yield and growth energy has demonstrated less successful. The growth rate of each organism eventually relies upon its intracellular concoction responses. Here we demonstrate that a thermodynamic model in light of a solitary, rate-restricting, enzymecatalysed response precisely depicts population growth of unicellular and multicellular living beings. All things considered these speak to every one of the three spaces of life, running from psychrophilic to hyperthermophiles, and including the highest temperature so far watched for growth ( $122^{\circ}\text{C}$ ). The outcomes give believable appraisals of thermodynamic properties of proteins and acquire, absolutely from life form natural growth rate information, connections between parameters previously recognized tentatively, therefore crossing over a hole between biochemistry and entire organism biology. We find that growth rates of both unicellular and multicellular living things can be depicted by the same temperature dependence model. The model results give solid help to a solitary exceedingly moderated response exhibit in the last universal common ancestor (LUCA). This is striking in that it implies that the growth rate reliance on temperature of unicellular and multicellular life shapes that developed over geological time ranges can be clarified by a similar model. Further research around there is required to develop models that would be valuable in process design and improvement.

**Keywords:** thermodynamic modeling, biological growth, successful, temperatures, population, develop models, process

### Introduction

Temperature governs the rate of chemical reactions including those enzymic forms controlling the development of life on Earth from singular cells to complex populations and traversing temperatures from well underneath solidifying to over the breaking point of water <sup>[1]</sup>. The growth rates of unicellular and multicellular organisms rely upon various procedures and steps, however all are on a basic level restricted by enzymic responses <sup>[2]</sup>. This acknowledgment gives a connection that conquers any hindrance between biochemistry and entire organism biology. By utilizing the suspicion of a solitary rate constraining response step we demonstrate that we can depict the growth rate of various poikilothermic living things. The temperature subordinate development bends of poikilothermic living beings over their biocenotic ranges have a trademark shape that may show up externally to be U-molded; however mindful examination demonstrates them to be more perplexing. The historical backdrop of previous approaches to depicting these bends is broad <sup>[3-6]</sup>. We utilize a model to describe the impact of temperature on biological systems that expect a solitary, rate-constraining, compound catalyzed response utilizing an Arrhenius shape that additionally takes into consideration protein denaturation. The relative accomplishment of microbial strains inside populations has been appeared to be fundamentally subject to protein denaturation <sup>[7]</sup>. Prior we introduced such a model and fitted it to 95 strains of organisms <sup>[8]</sup>. In this work notwithstanding information on microorganisms, we likewise incorporate information on the

intrinsic growth rates for bugs and acari got from lifetable analysis and find that these multicellular strains are additionally very much depicted by the model. Eminent among the displayed strains is the consideration of hyperthermophiles dynamic at the most elevated temperatures so far known for biological growth ( $121^{\circ}\text{C}$  <sup>[9]</sup>,  $122^{\circ}\text{C}$  <sup>[10]</sup>). The most minimal temperature modeled was  $22^{\circ}\text{C}$ , beneath which growth rates can't be dependably contrasted due with ice arrangement and the zone of warm capture. In this paper we address biological implications and results emerging from examination of significantly more broad information than beforehand utilized <sup>[8]</sup> and by gathering strains by their thermal optima as opposed to by taxonomy.

### Review of Literature

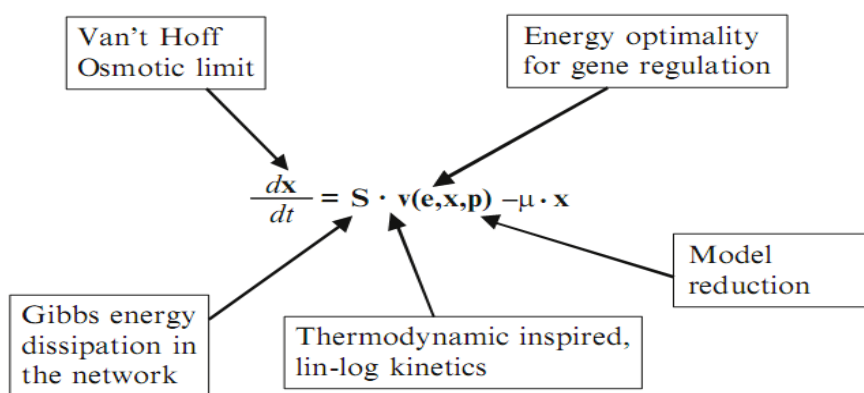
In this paper we build up the thermodynamic of biological cells growth, with specific respects to conceivable control of the cells growth by a control of the particles transport over the cell membrane. To do as such, we consider that cells unexpectedly trade warmth, and this warmth is identified with their biochemical and biophysical behaviour. This squandered warmth speaks to the cooperation between the cell and its condition, a kind of "spontaneous communication" towards condition. This collaboration is basic to creating a thermodynamic investigation of the cell. Without a doubt, cells are excessively mind boggling, making it impossible to comprehend the contribution of each process to the global result, and the investigation of cells as secret elements enables us to improve the examination by considering just the inflow

and outpouring adjusts. In addition, it is less demanding to approach the cell environment than to the living cell itself. These contemplations enable us to present the bases of the bio thermodynamic approach presented in the study of the cells:

1. An open irreversible real linear or non-linear system is considered;
2. Each process has a finite lifetime  $\tau$ ;
3. What happens in each instant in the range  $[0, \tau]$  cannot be known, but what has happened after time  $\tau$  (the result of the process) is well-known (at least it is sufficient to wait and observe): local equilibrium is not necessarily required;
4. The balance equations are balance of fluxes of energy, mass and ions.

Recently, it has been featured that any impact in Nature is dependably the result of the dynamic adjusts of the communications between the genuine frameworks and their environments <sup>[12]</sup> and the real systems advancement is constantly identified with the decline of their free vitality, at all time. Along these lines, bio thermodynamics is constructing just in light of two fundamental ideas of physics: interactions and streams. The outcome is the systematic definition of stream based investigation in thermodynamics, which can assume the part of an "encouraging point" of the diverse modelling approach to bio systems. To be sure, on the off chance that we consider regular frameworks we can feature that they are constantly open frameworks, which implies that they can trade warmth and mass with their environment. In this way, the collaboration with the earth is a fundamental concept for the thermodynamic analysis.

**Thermodynamic Principles in Mathematical Models of Biological growth Systems:** The key aspect of a living cell is



**Fig 1:** Impact of thermodynamic principles in systems biology

In the following sections we will show that thermodynamic principles can be used to shed light on this information (Fig. 1):

- Metabolite concentration levels ( $X_j$ ) and their control mechanisms
- The stoichiometry of growth ( $S$ )
- The genetic regulation of enzyme levels ( $e_i$ )
- Principles of model reduction
- The kinetics of enzyme catalyzed reactions based on thermodynamic driving force.

the arrangement of new cells, called growth. Growth requires that a cell produce each of the particles display in the recently formed cells. This happens in a vast and complex metabolic (response) network. This network is made out of numerous responses, which expend and create small molecules, called metabolites. Each reaction is catalyzed by a particular protein, which is under genetic/environmental control. Expectation of growth requires a mathematical model of this system for which the fundamental equations are the mass adjusts of intracellular metabolites. In vector notation:

$$\frac{dX}{dt} = SV(e, X, p) - \mu X, \tag{1}$$

Where  $X$  is the vector containing the individual intracellular metabolites  $X_j$ .  $V$  is the vector of the rates of enzyme catalyses reactions, with  $v_i$  the rate of reaction catalyzed by enzyme present at an amount  $e_i$ . The rate  $v_i$  depends on the amount of enzyme present,  $e_i$ , on the kinetic effect of metabolites  $X_j$  involved (e.g. substrate, product and possible allosteric effectors) and the parameters  $p$  (e.g.  $V^{max}$ , affinities, Hill coefficient etc.).  $S$  is the so-called stoichiometric matrix which represents the structure of the reaction network. Its rows represent the metabolites, its columns the reactions. The term  $\mu X$  is the so-called dilution term.

Equation (1) requires information on the dynamic behaviour and values of metabolite concentrations, on the values of stoichiometric coefficients, on enzyme levels resulting from genetic regulation and on the shape/algebraic nature of the enzyme kinetic relations, and (1) is the basis of parameter estimation from experimental data and the associated need for model reduction.

**Thermodynamic relationships:** The probability of the local (chemically dynamic) state for the thermal groups we allude to the last as local state bends <sup>[8]</sup> since they speak to the extent of the rate-controlling compound that is in the local compliance. The bends for the likelihood of the local state have brought down tops for psychrophilic, mesophilic, and Ascomycota, and the bends are taller and continuously leveled for thermophiles and hyperthermophiles. The higher and compliment crest for the thermophiles and hyperthermophiles suggests protein strength over an undeniably expanded

temperature range. The lower top levels for the lower temperature groups may be translated as lessened soundness for psychrophilic <sup>[10]</sup> and Ascomycota proteins <sup>[9]</sup>. The psychrophilic local state bend is likewise moved to one side of alternate gatherings, which are all approximately aligned at the same lower temperature (&275K). The deviation of the psychrophilic below alternate gatherings proposes that an unthinking contrast has developed separating psychrophilic from the other groups.

**Bio thermodynamics of biological cells:** Cells are open complex thermodynamic systems. They can be additionally viewed as mind boggling motors that execute a progression of chemical reactions. Energy transformations, thermo-electro-synthetic procedures and transports phenomena can happen over the cells membranes. Moreover, cells can likewise effectively adjust their practices in connection to changes in their environment. Nature, from a physical, biological, chemical and mathematical point of view, is a perplexing framework, while from perspective; it is the "primary" designer! Specifically, cells can be displayed as versatile warm and compound motors which change over vitality in one shape to another by coupling metabolic and chemical reactions with transport processes <sup>[11]</sup>, by devouring irreversibly free vitality for warm and chemical processes, transport of issue, vitality and particles.

Energy is a thermodynamic property of any framework in connection to a reference state, which changes amid any procedure, while its aggregate sum stays steady in connection to the universe, considering it as the framework together with its environment. In cells, many processes such as replication, interpretation and interpretation need to change over sub-atomic binding energy, chemical bond hydrolysis and electromagnetic gradients into mechanical work, identified with conformational changes and relocations <sup>[9]</sup>. The biomechanical analysis of DNA has called attention to the associations among powers, thermodynamics, nano-mechanical and electromagnetic conduct of biological structures and energy <sup>[10]</sup>. Thermodynamics is the science which studies both vitality and its best use in connection to the accessible vitality assets with specific respects to energy conversion, including power generation, refrigeration and connections among the properties of issue, including likewise living issue. In this way, thermodynamics can be presented in the mechanobiological and system biological approach so as to enhance these sciences by investigating the bio frameworks additionally from a thermal point of view: a new bioscience could be viewed as, the bio thermodynamics. To be sure, the first law of thermodynamics expresses the protection of vitality, while the second law states that entropy constantly increments for the framework and its environment and presents the measurable and informational meaning of global quantities <sup>[11]</sup>.

**Thermodynamics of population growth:** Reaction diffusion models have been utilized to portray different phenomena in liquid flow, dendritic growth, population growth, beat engendering in nerves, and other biological phenomena. The following conditions are gotten from the established dispersion condition considering a source term of particles;

the most generally one utilized as a part of the investigation of population growth is the Fisher condition, in which the source term is logistic <sup>[1]</sup>. Some authors have summed up this investigation by presenting dispersion coefficients relying upon the number thickness, so that the problem becomes highly nonlinear. Notwithstanding, no memory is considered in the latter models.

In the present paper we sum up the past response diffusion models in biological populations by including memory effects. It is notable that an animal's motion amid a little era tends to continue an indistinguishable way from it did in the quick time frame some time recently. This memory has as an immediate consequence the postponement in the presence of the population flux, a defer that has not been considered in past models as far as anyone is concerned. Give us a chance to see, then again, that postponed transport equations have been generally utilized to solve diffusion problems, hyperbolic warmth conduction, and gooey transport processes. These sorts of transport equations demonstrate an important feature.

**The nature of the rate-limiting reaction:** While the model performs astoundingly, both regarding its general consistency with protein biochemistry and in the great fits acquired, some predictions don't completely concur with thermodynamic expectations and there exists the likelihood that the basic component might be more complex than a solitary, rate-constraining, catalyst catalyzed response. By and by, the model underlines the significance of thermodynamics in biological processes particularly those identifying with the cooperation between proteins and water particles, which thusly may rely upon the properties of water itself. Be that as it may, on the off chance that it takes the type of a single reaction then we can estimate on its nature. An instrument by which cells control denaturation might be proposed by thought of protein chaperones. Some cases are the bacterial chaperonins. Such frameworks act amid all over again collapsing and to refold unfurled substrate proteins. They are activated by the swelled presentation of hydrophobic gatherings in the unfurled proteins. GroEL and GroES function together to make an Anfinsen hydrophilic confine containing charged deposits that aggregate requested water atoms, making the substrate protein cover its hydrophobic buildups and refold into its local state. The rate at which the GroEL and GroES work continues is controlled by ATP hydrolysis. In the event that warmth shock proteins speak to the rate-restricting stride, the rate at which they work must be the basic factor. Those chaperones that are in charge of once more collapsing and refolding are ATP-subordinate. Articulation of important chaperones (GroEL, GroEL, GrpE, DnaK) appear to end up noticeably noiseless as bacterial cells kick the bucket from sudden thermal stress <sup>[59]</sup>. Along these lines, we theorize that the rate-constraining might be connected to a process leading to or specifically connected to protein folding. Responses conceivably incorporate a scope of important enzymes either ordering or supporting protein collapsing with denaturation of the response prompting hindrance of the broad protein folding process. Conceivable illustrations incorporate trigger factor, peptidylprolyl

isomerases (the moderate stride in protein folding), protein disaggregation and support of ATP accessibility to the folding system.

### Conclusion

Life is an authoritative and thermodynamic process that tends towards the maximum conversion of available energy. The biochemical reactions create or expend external metabolites, and they associate internal metabolites, in consistent fixations in the cells at their unfaltering states. To do as such, the cell must trade vitality and matter through its film. The fundamental phenomena utilized by cells to achieve their optimality comprise of a redistributing of the flow patterns through their metabolic network. By utilizing the bio thermodynamics, it has been featured how the different ions have different effect on the utilization of vitality by the cell to grow. To do as such, a control of the cells behaviours is presented. Here, an electromagnetic field is utilized as a control system, yet other field could be utilized. Cells inside and outside an electromagnetic field have been considered. The positive ions determine a diminishing of the energy utilized by the disease, with the end goal that the disease can't develop as outside the field. Then again, the negative particle builds the utilization of energy. It implies that a control of ions can decide a control of the volume growth of a disease. This outcome can be stretched out to all the atomic fluxed over the cell membrane, acquiring a conceivable bio thermodynamic way to deal with control the disease growth. Thermodynamic principles do shape the properties of biological systems, with extensive and very fascinating results for their required in systems biology.

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