

Short Note

A Simple Thermodynamic Analysis of Photosynthesis

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Abstract: In this paper we present a comparative study of nine photosynthetic pathways by means of their thermodynamic performance. The comparison is made by using the thermal efficiency of light-to-chemical energy conversion and the so-called ecological criterion arising from finite-time thermodynamics. The application of both criteria leads to photosynthesis made by metaphytes and non sulfur purple bacteria as those of best thermodynamic performance. In spite of the simplicity of our thermodynamic approach some insights over the low overall efficiency of photosynthesis is suggested.

Keywords: photosynthesis, thermodynamic performance, ecological function.

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1. Introduction

Schrödinger suggested that the maintenance of high organization of living beings is due to a continuum influx of negative entropy [1]. Photosynthesis is a process where energy-rich organic molecules emerge from simple, energy-poor molecules absorbing solar photons [2]. This photochemical reaction occurs in the photosynthetic reaction center, which is a very complicated molecular complex [3]. Many models to describe the photosynthetic center have been proposed. Van Rotterdam et. al. [3] suggested that the transduction of photons' energy to a transmembrane electrochemical potential difference for protons operates in a simple battery-like manner. De Vos [4],

conceived the photosynthesis engine as composed by two parts: a photovoltaic component that absorbs the solar radiation and converts it into work, and a chemical reactor, which uses the work in order to keep a chemical reaction going on in the "reverse direction". A very complete model was proposed by Juretic and Zupanovic [5]. This model is based on a non-equilibrium thermodynamics approach provided by Meszena and Westerhoff [6]. In the treatment of photosynthesis several design principles for biological systems have been used, such as maximum efficiency [7] & [8], minimum entropy production [2] and maximum entropy production [5]. Recently Lavergne [9] has disscused a photochemical energy transducer as a model for photosynthesis within a second law analysis. Nowadays, there is no consensus about what optimization criterion if any is followed by photosynthesis performance. In this article we present a brief and simple comparison between nine photosynthetic pathways in terms of their overall thermal efficiencies and also in terms of the so-called ecological function defined within the context of finite-time thermodynamics [10]. By means of both criteria we found that photosynthesis occurring in superior plants and non sulfur purple bacteria has a better performance than the other ones. In fact, in our "ecological" analysis, we use the integral of the ecological function over the duration of photosynthetic chemical reaction. Although, efficiency and the integral of the ecological function have a similar dependence on the free energy changes, we show both comparisons because the integral of ecological function exhibits some features that are not present in the efficiency behavior. The article is organized as follows: In sect. II we present the basis and main assumptions of our thermodynamic analysis in a first approximation; in sect. III we show our thermal efficiency calculations; in sect. IV the analysis based on the ecological function is shown; in sect. V we present again efficiency and ecological calculations taking into account the role of dilution of Sun's radiation. Finally, in sect. VI, we present the concluding remarks.

2. Photosynthesis Thermodynamics

There exist several photosynthetic pathways by means of which living organisms can store solar energy in form of chemical energy. The most studied pathway being the photosynthesis made by superior plants and cyanobacteria, which can be summarized as follows,

$$6CO_2 + 6H_2O \longrightarrow C_6H_{12}O_6 + 6O_2$$

However, alternative pathways are used by green and purple bacteria, which use compounds different from water, as sources of hydrogen to synthesize glucose. Among these compounds donors of hydrogen are sulfhydric acid, and several organic compounds such as methanol and ethanol for example. In Appendix we show nine photosynthetic pathways with their corresponding standard free energy changes. The photosynthesis reaction is usually divided in two groups of chemical reactions. The first one called the light phase of photosynthesis, in which the reactions are driven by solar light and the second one, called the dark phase, where the chemical reactions occur without solar light presence [11-15]. All chemical steps of photosynthesis in superior plants are well described in [11-15]. In the following sections we will develop a thermodynamic analysis of photosynthesis within the context of some results arising from classical and finite-time thermodynamics.

Our thermodynamic analysis of photosynthesis starts by establishing the following convenient working hypothesis:

- a) The Sun, the Earth and the photosynthetic organism (PO) are three different thermodynamic systems.
- b) The Sun is a thermal reservoir with constant temperature $T_s = 5762$ K [4].
- c) The Sun has constant pressure, volume and chemical composition.
- d) Earth behaves as a thermal reservoir at $T_E = 298.15$ K.
- e) The Earth is a system with constant pressure, volume and chemical composition.
- f) The photosynthetic organism (PO) has constant pressure, volume and temperature, with $T_{PO} = T_E = 298.15$ K.
- g) The PO chemical composition is not constant.
- h) All photosynthetic reactions are isothermal processes at $T_{PO} = 298.15$ K.

For our thermodynamic study we will divide the overall process in three steps: i) The light travels from de Sun up to the PO without making any work; ii) The PO uses some part of the received energy to produce glucose by using some chemical compounds; iii) The PO delivers the remaining energy to the Earth in form of heat (see Figure 1).



Figure 1. Diagram corresponding to the overall energy fluxes.

Consider the following thermodynamic equations [16]:

$$dU = TdS - PdV + \sum_{k=1}^{N} \mu_k dn_k \qquad (1)$$
$$dG = -SdT + VdP + \sum_{k=1}^{N} \mu_k dn_k \quad , \qquad (2)$$

where U is the internal energy, G the Gibbs free energy, T the temperature, P the pressure, V the volume, S the entropy, μ_k the *k*-th chemical potential and n_k is the *k*-th number of moles. From Eqs. (1) and (2) and the mentioned working hypothesis we get the following expressions:

For the Sun system we have:

$$dU_s = T_s dS_s \qquad (3a)$$

$$dG_s = 0 \quad , \qquad (3b)$$

the subscript "S" refers the Sun.

For the Earth system, we obtain:

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$$dU_E = T_E dS_E \qquad (4a)$$

$$dG_E = 0 \qquad , \qquad (4b)$$

the subscript "*E*" refers the Earth.

Finally, for the PO we get:

$$dU_{PO} = T_{PO} dS_{PO} + \sum_{k=1}^{N} \mu_k dn_k$$
 (5a)

$$dG_{PO} = \sum_{k=1}^{N} \mu_k dn_k \qquad , \qquad (5b)$$

the subscript "PO" refers the photosynthetic organism.

From Eqs. 5a and 5b, we can see that the chemical work is directly the Gibbs free energy change and thus:

$$dU_{PO} = T_{PO} dS_{PO} + dG_{PO} \quad . \tag{6}$$

By integrating Eqs. 3a, 4a, and 6 we obtain:

$$\Delta U_s = T_s \Delta S_s \tag{7}$$

$$\Delta U_E = T_E \Delta S_E \tag{8}$$

$$\Delta U_{PO} = T_{PO} \Delta S_{PO} + \Delta G_{PO} . \tag{9}$$

Thermodynamics of Step #1

Several authors [2] assert that superior plants need 60 photons to synthesize one glucose molecule. Then for each glucose mol synthesized, the Sun losses energy given by:

$$\Delta U_{S}^{Step\#1} = -\frac{60N_{A}hc}{\lambda} \quad , \qquad (10)$$

being N_A the Avogadro's number, *h* the Planck's constant, *c* the light's speed and λ the photon's wavelength. With this energy change there is a concomitant entropy change, expressed as:

$$\Delta S_s^{Step \# 1} = -\frac{60N_A hc}{\lambda T_S}.$$
 (11)

The energy lost by the Sun is absorbed by the photosynthetic organism, then:

$$\Delta U_{PO}^{Step\#1} = \frac{60N_A hc}{\lambda} \quad , \qquad (12)$$

and therefore, with an entropy gain given by:

$$\Delta S_{PO}^{Step\#1} = \frac{60N_A hc}{\lambda T_P} . \qquad (13)$$

Earth does not participate in this step of the process, therefore,

$$\Delta U_E^{Step\#1} = 0$$
 and $\Delta S_E^{Step\#1} = 0$,

thus, the total change of entropy in this step is:

$$\Delta S_{Total}^{Step \# 1} = \frac{60N_A hc}{\lambda} \left(\frac{1}{T_{PO}} - \frac{1}{T_S} \right) \quad . \tag{14}$$

To obtain Eq. (14) we followed the first approximation used by Brittin and Gamow [17]. This approach consists in assuming that the diluted radiation stemming from Sun reaches the Earth with a grey body radiation spectrum. Then the process taking place when energy exchange between different frequencies is permitted can be compared to a flow of heat from a reservoir at the temperature T_S to a cooler reservoir at the temperature T_E . (See figure 1. of [17]).

Thermodynamics of Step #2

In this step of the process the PO uses part of the absorbed energy in Step#1 and transforms it as chemical energy in the glucose synthesis. In this step, the PO does not exchange energy with its sorroundings, then $\Delta U_{PO}^{\text{Step#2}} = 0$ and the entropy change turns out to be:

$$\Delta S_{PO}^{Step#2} = -\frac{\Delta G_{PO}^{Step#2}}{T_{PO}}, \quad (15)$$

being $\Delta G_{PO}^{\text{Step#2}}$ the PO free energy change in Step#2.

Since Earth and Sun do not participate in this step both their entropy and energy changes become zero. Thus, the total entropy change in Step#2 becomes:

$$\Delta S_{Total}^{Step#2} = -\frac{\Delta G_{PO}^{Step#2}}{T_{PO}} .$$
 (16)

Thermodynamics of Step#3

During this step the PO rejects to Earth the remaining energy, which was not used in glucose synthesis. Thus the internal energy and entropy changes for the PO are:

$$\Delta U_{PO}^{Step#3} = -\left(\Delta U_{PO}^{Step#1} - \Delta G_{PO}^{Step#2}\right)$$
(17)
$$\Delta S_{PO}^{Step#3} = -\frac{\left(\Delta U_{PO}^{Step#1} - \Delta G_{PO}^{Step#2}\right)}{T_{PO}} ,$$
(18)

and for the Earth these changes are:

$$\Delta U_E^{Step#3} = \left(\Delta U_{PO}^{Step#1} - \Delta G_{PO}^{Step#2} \right)$$
(19)
$$\Delta S_E^{Step#3} = \frac{\left(\Delta U_{PO}^{Step#1} - \Delta G_{PO}^{Step#2} \right)}{T_E}.$$
(20)

Finally, for the Sun we have $\Delta U_S^{\text{Step#3}} = 0$ and $\Delta S_S^{\text{Step#3}} = 0$.

Thus, by adding Eqs. (18) and (20) with $\Delta S_s^{\text{Step#3}} = 0$, the total entropy change in Step#3 is:

$$\Delta S_{Total}^{Step \# 3} = 0 \quad , \tag{21}$$

and therefore, by using Eqs. (14), (16) and (21), the total entropy change in the three steps is:

$$\Delta S_{Universe} = \frac{60N_A hc}{\lambda} \left(\frac{1}{T_{PO}} - \frac{1}{T_S} \right) - \frac{\Delta G_{PO}^{Step#2}}{T_{PO}}$$
(22)

This expression can be simplified if we take into account the Brittin and Gamow's approximation [17]:

$$\frac{1}{T_s} \approx 0 \ .$$

Then Eq. (22) becomes:

$$\Delta S_{Universe} \approx \frac{60N_A hc}{\lambda T_{PO}} - \frac{\Delta G_{PO}^{Step#2}}{T_{PO}}, \quad (23)$$

3. Efficiency Calculation

In this section we compare the efficiency of the nine photosynthetic pathways shown in the Appendix. Here, the efficiency is taken as:

$$\eta(\Delta G^0, \lambda) \equiv \frac{\Delta G^0}{60N_A hc / \lambda} \qquad (24)$$

Where ΔG^0 is the standard Gibbs free energy change during the photosynthetical reaction. The ΔG^0 and λ data for each of the nine cases are shown in the Appendix. The numerical results obtained from Eqs. (23) and (24) are depicted in Table I. The wavelength values were taken from [13], where it is asserted that the light used in photosynthesis by some PO's have the maxima bounds shown at the second column of Table I. In Eq. (23) and (24) we use the standard free energy because of the difficulty to obtain actual free energy changes. (In addition, we are more interested in comparing efficiency values than in their absolute values). This issue has been discussed by Cornish-Bowden [18], but some authors [12] have used standard Gibbs free energy changes in efficiency calculations.

Table 1. Efficiency Eq. (24) and entropy change Eq. (23) for the nine photosynthetic pathways given in the Appendix. For several pathways different λ values were taken. Clearly, reaction 1 (superior plants) has the higher efficiency.

Reaction	λ [nm]	ΔU [kJ/mol]	ΔG^0 [kJ/mol]	η %	ΔS _{Universe} [kJ/(mol K)]
1	680	10555 287	2880.31	27 288	25 742
1	840	10555.207 9544 756	420.51	5 029	23.742
2	840	8344./30	429.04	5.028	27.218
2	870	8250.109	429.64	5.208	26.230
2	890	8064.713	429.64	5.327	25.608
3	870	8250.109	744.57	9.025	25.174
3	890	8064.713	744.57	9.232	24.552
4	840	8544.756	621.47	7.273	26.575
4	870	8250.109	621.47	7.533	25.587
4	890	8064.713	621.47	7.706	24.965
5	840	8544.756	584.86	6.845	26.698
5	870	8250.109	584.86	7.089	25.709
5	960	7476.661	584.86	7.822	23.115
6	840	8544.756	71.27	0.834	28.420
6	870	8250.109	71.27	0.864	27.432
6	960	7476.661	71.27	0.953	24.838
7	870	8250.109	1066.56	12.928	24.094
7	960	7476.661	1066.56	14.265	21.500
8	870	8250.109	609.48	7.388	25.627
8	960	7476.661	609.48	8.152	23.033
9	798	8994.480	320.65	3.565	29.092

4. Comparison of the Photosynthesis Ecological Functions

In this section we will present a brief analysis of some photosynthetical reactions within the context of finite-time thermodynamics [2], [5-8], [10]. In particular, we will use the so-called ecological criterion [10]. This criterion is based in the maximization of the so-named ecological function, defined as:

$$E = P - T\sigma , \qquad (25)$$

where P is the power output of an energy conversion process, T is the absolute temperature of a heat reservoir to which the heat is rejected and σ is the total entropy production during the process.

To obtain an expression for *E* independent of time, we integrate Eq. (25) along the time interval τ employed to synthesize one mol of glucose by using 60 mol of photons, that is:

$$\int_0^\tau E dt = \int_0^\tau (P - T\sigma) dt \quad . \tag{26}$$

We take this process as an isothermal one that is, $T=T_{PO}$, then:

$$\int_0^\tau E dt = W - T \Delta S_U \quad , \tag{27}$$

where W is the mean work done, and ΔS_U is the mean total entropy change of the universe, both during the time interval τ . This interval can be different for each photosynthetic pathway, due to they are distinct in many respects, including structural, kinetic and thermodynamic aspects. However, we will only compare the integral of the ecological function, see 6th column of Table II.

On the other hand, by means of Eq. (23) we have:

$$\Delta S_U \approx \frac{1}{T_{PO}} \left(\Delta U_{PO}^{Step \# 1} - \Delta G_{PO}^{Step \# 2} \right), \qquad (23)$$

with:

$$\Delta U_{PO}^{Step\#1} = \frac{60N_A hc}{\lambda} , \qquad (12)$$

and we also have:

$$W = \Delta G_{PO}^{Step \# 2} . \tag{28}$$

Thus, the integral of ecological function is:

$$\int_{0}^{\tau} E dt = \Delta G_{PO}^{Step \# 2} - T_{PO} \cdot \frac{1}{T_{PO}} \left(\Delta U_{PO}^{Step \# 1} - \Delta G_{PO}^{Step \# 2} \right)$$
$$\int_{0}^{\tau} E dt = 2\Delta G_{PO}^{Step \# 2} - \Delta U_{PO}^{Step \# 1}, \qquad (29)$$

that is a function of ΔG and λ , $\int_0^\tau E dt = f(\Delta G, \lambda)$, given by,

$$\int_{0}^{\tau} E dt = 2\Delta G_{PO}^{Step\#2} - \frac{60N_{A}hc}{\lambda}$$
(30)

We compare now the photosynthetical pathways by means of the values of the integral of ecological function. In Table II, we show the results of Eq. (30) for nine photosyntetical pathways using the standard Gibbs free energy.

As we can see, the photosynthesis in superior plants has the maximum value for the integral of the ecological function. Here, we only compare the integral of the ecological function for the nine photosynthetical pathways searching for its maximum value and we had not made a general optimization of the ecological function. It has been suggested that the ecological function represents a good trade off between high power output and low entropy production [10], [19]. Thus, the photosynthesis made by metaphytes is better than the other ones, in the sense of the ecological function. Remarkably, in all of the calculated cases, the integral of function *E* has negative values. According with Eq. (25), this means that the dissipative term $T\sigma$ is larger than the power term *P*, which may be a possible explanation of the low overall efficiency of photosynthesis [20]. Although efficiency and $\int_0^r Edt$ have a similar dependence on ΔG^0 , their results are not redundant because the integral of the ecological function gives information about the dissipative terms that are not explicitly taken into account in the efficiency analysis. In addition, the integral of ecological function and the efficiency have dependence on the wavelength, and if we simultaneously analyze the dependence on ΔG^0 and λ it is easy to see that both functions have not exactly the same monotony (see Figure 2).

5. Analyzing the Radiation's Entropy Change.

Such as Brittin and Gamow assert [17], Eq. (14) is only an approximation. However, if a more complete estimation of $\Delta S_{Universe}$ for its radiative term is made, we must calculate the correct entropy change for the radiation during its travel from the Sun up to the Earth. Then, it is necessary to do a new thermodynamic analysis considering four steps during the overall process: i) The radiation travels from de Sun up to the Earth suffering a free adiabatic expansion without making any work, and without exchanging any heat; ii) The PO absorbs light from the environment; iii) The PO uses some part of the received energy to produce glucose by using some chemical compounds; iv) The PO delivers the remaining energy to the environment (see Figure 3). We now give a brief thermodynamic description of the four steps.

Depation	2 [nm]	AU [], I/mol]	AC ⁰ []r I/mol]	m 9/	C T
Reaction	∧ [nm]	ΔU [KJ/moi]	ΔG [KJ/III01]	η %	$\int_0^{\infty} E dt \ [kJ/mol]$
1	680	10555.287	2880.31	27.288	-4794.667
2	840	8544.756	429.64	5.028	-7685.476
2	870	8250.109	429.64	5.208	-7390.829
2	890	8064.713	429.64	5.327	-7205.433
3	870	8250.109	744.57	9.025	-6760.969
3	890	8064.713	744.57	9.232	-6575.573
4	840	8544.756	621.47	7.273	-7301.816
4	870	8250.109	621.47	7.533	-7007.169
4	890	8064.713	621.47	7.706	-6821.773
5	840	8544.756	584.86	6.845	-7375.036
5	870	8250.109	584.86	7.089	-7080.389
5	960	7476.661	584.86	7.822	-6306.941
6	840	8544.756	71.27	0.834	-8402.216
6	870	8250.109	71.27	0.864	-8107.569
6	960	7476.661	71.27	0.953	-7334.121
7	870	8250.109	1066.56	12.928	-6116.989
7	960	7476.661	1066.56	14.265	-5343.541
8	870	8250.109	609.48	7.388	-7031.149
8	960	7476.661	609.48	8.152	-6257.701
9	798	8994.480	320.65	3.565	-8353.180

Table 2. Comparison of nine photosynthetical pathways in terms of Ecological Function given by Eq. (30). For this case, reaction 1 (superior plants) has the higher value of the integral of the ecological function.



Efficiency %

Figure 2. The integral of the ecological function in terms of the efficiency for each ΔG^0 of the nine pathways. Data obtained by using Eqs. (24) and (30).



Figure 3. Alternative diagram corresponding to the overall energy fluxes.

Thermodynamics of Step #1

As Figure 3 shows, the photons in the surface of the Sun act as a black-body radiation at $T_S = 5762$ K that suffers a free adiabatic expansion during the travel across the space. Just before arriving to the Earth, the diluted gas of photons has the properties of an equivalent grey-body at $T_S = 5762$ K in a radiative equilibrium with the Earth's atmosphere which acts as a black-body at $T_E = 298.15$ K [17]. The emissivity ε of this equivalent grey-body radiation is the square of the Sun's radius divided by the square of the radius of the Earth's orbit which is the so-called dilution factor [4]. Then the radiation equilibrium equation is:

$$\varepsilon \cdot \sigma \cdot T_s^4 = \sigma \cdot T_E^4 \quad , \tag{31}$$

$$\varepsilon = \frac{R_s^2}{R_o^2} , \qquad (32)$$

where R_S is the radius of the Sun, R_O is the radius of the Earth's orbit and σ is the Stefan-Boltzman's constant. The initial state of the photons' gas is the black-body radiation at $T_S = 5762$ K in the Sun's surface having entropy S_S , energy U_S and volume V_S . On the other hand, the final state is the black-body radiation at $T_E = 298.15$ K in the Earth's atmosphere having entropy S_E , energy U_E and volume V_E . Then the entropy and the energy of the radiation at the initial and final states are given by [21]:

S_{s}	$=\frac{4}{3} \cdot a$	$\cdot T_S^3 \cdot$	V_{S}	,	(33)
	5	5			· · ·

$$S_E = \frac{4}{3} \cdot a \cdot T_E^3 \cdot V_E \quad , \tag{34}$$

$$U_s = a \cdot T_s^4 \cdot V_s \quad , \tag{35}$$

$$U_E = a \cdot T_E^4 \cdot V_E \quad , \tag{36}$$

where *a* is defined by $a = 4\sigma / c$.

Due to energy's conservation, we must consider $U_S = U_E = 60N_Ahc / \lambda$ and because of the geometric effect of the dilution we have $\varepsilon = V_S / V_E$ [4], [17]. By using these facts along with the Eqs. (33), (34), (35) and (36) we get the Brittin and Gamow's formula for the radiation's entropy change [17]:

$$\Delta S_{Total}^{Step \# 1} = \frac{4}{3} \left(\frac{60N_A hc}{\lambda} \right) \left(\frac{1}{T_E} - \frac{1}{T_S} \right) .$$
(37)

with

Thermodynamics of Step #2

During this step, the light energy goes from the Earth's atmosphere at $T_E = 298.15$ K to the PO at the same temperature $T_{PO} = 298.15$ K. Then the entropy change becomes zero.

$$\Delta S_{Total}^{Step#2} = 0 . aga{38}$$

Thermodynamics of Step #3

During this step, the PO uses some energy in the photosynthetic reaction to making chemical work denoted by $\Delta G_{PO}^{\text{Step#3}}$. Then the entropy change during this step is:

$$\Delta S_{Total}^{Step#3} = -\frac{\Delta G_{PO}^{Step#3}}{T_{PO}} . \tag{39}$$

Thermodynamics of Step #4

Finally, the PO at $T_{PO} = 298.15$ K delivers the remaining energy to the environment at the same temperature $T_E = 298.15$ K. Then the entropy change becomes zero.

$$\Delta S_{Total}^{Step#4} = 0 \quad . \tag{40}$$

By using Eqs. (37), (38), (39) and (40) and simplifying by means of the approximation $1/T_s \approx 0$, we obtain:

$$\Delta S_{Universe} \approx \frac{4}{3} \left(\frac{60N_A hc}{\lambda T_{PO}} \right) - \frac{\Delta G_{PO}^{Step \# 3}}{T_{PO}} , \qquad (41)$$

and now by taking Eqs. (27) and (41) we get:

$$\int_{0}^{\tau} E dt = 2\Delta G_{PO}^{Step\#3} - \frac{4}{3} \left(\frac{60N_{A}hc}{\lambda} \right)$$
(42)

Finally, by using Eqs. (24), (41) and (42) in the comparison of the nine photosynthetical pathways mentioned, we arrive to Table III and Figure 4. As it can be seen, under this analysis the maximum value of $\int_0^{\tau} Edt$ is now reached by the photosynthesis made by non sulfur purple bacteria. (Reaction number seven from the appendix with its superior value $\lambda = 960$ nm [13].)

			-			
Reaction	λ [nm]	ΔU [kJ/mol]	∆G ⁰ [kJ/mol]	η %	ΔS _{Universe} [kJ/(mol K)]	$\int_0^\tau E dt \ [kJ/mol]$
1	680	10555 287	2880.31	27 288	27 5/2	8312.006
1	060	10555.267	2000.51	27.200	57.545	-8313.090
2	840	8544./56	429.64	5.028	36.//1	-10533.728
2	870	8250.109	429.64	5.208	35.454	-10140.865
2	890	8064.713	429.64	5.327	34.625	-9893.671
3	870	8250.109	744.57	9.025	34.397	-9511.005
3	890	8064.713	744.57	9.232	33.568	-9263.811
4	840	8544.756	621.47	7.273	36.128	-10150.068
4	870	8250.109	621.47	7.533	34.810	-9757.205
4	890	8064.713	621.47	7.706	33.981	-9510.011
5	840	8544.756	584.86	6.845	36.251	-10223.288
5	870	8250.109	584.86	7.089	34.933	-9830.425
5	960	7476.661	584.86	7.822	31.474	-8799.161
6	840	8544.756	71.27	0.834	37.973	-11250.468
6	870	8250.109	71.27	0.864	36.656	-10857.605
6	960	7476.661	71.27	0.953	33.197	-9826.341
7	870	8250.109	1066.56	12.928	33.317	-8867.025
7	960	7476.661	1066.56	14.265	29.859	-7835.761
8	870	8250.109	609.48	7.388	34.850	-9781.185
8	960	7476.661	609.48	8.152	31.392	-8749.921
9	798	8994.480	320.65	3.565	39.148	-11351.340

Table 3. Comparison of nine photosynthetical pathways in terms of Ecological Function given by Eq. (42). In this case, the higher value of the integral of the ecological function is reached by reaction 7 with $\lambda = 960$ nm.



Integral of the Ecological Function vs. Efficiency

Figure 4. The Integral of the Ecological Function in terms of the Efficiency. Upper plot corresponds Eqs. (24) and (30). Inferior graph corresponds to Eqs. (24) and (42).

6. Concluding Remarks

In many man-made energy converters several design principles have been used to obtain certain optimal objectives [22-26]. Among these, one finds maximum power output, minimum entropy generation, maximum efficiency and several others. However, is a pertinent hypothesis to think that Nature uses analogous design procedures? Nowadays, there is not a consensus about this matter. However, several authors have proposed optimization criteria to analyze the performance mode of operation of some natural processes. In particular, for the case of photosynthesis, criteria such as maximum efficiency, minimum entropy production and maximum entropy production have been used. We have described a simplified analysis of photosynthesis, in particular, we compared the thermal efficiency of light-to-chemical energy conversion for nine different photosynthetic pathways. Additionally, we also compared these chemical pathways by means of the so-called ecological function used in finite-time thermodynamics. These analyses are not redundant because the "ecological" criterion reveals some features that are not explicit in the efficiency analysis. Under this analysis, in Table I we see that reaction 1 (metaphytes) reaches the higher overall thermal efficiency. Under the first Brittin and Gamow's approach, the ecological analysis gave that metaphytes reach the higher value of the integral of the ecological function. However, when the second Brittin and Gamow's approach is taken into account (the 4/3 term), the non sulfur purple bacteria slightly reach the higher value of the integral of the ecological function above the corresponding value for the metaphytes. It is convenient to remark that our calculation for the thermal efficiency must be taken as an inferior bound of actual efficiencies due to typical standard Gibbs free energy changes are smaller than actual Gibbs free energy changes [12]. In spite of our analysis is based on a very elementary approach, it suggests that possibly some biological processes are concomitant with certain thermodynamic optimality criteria. In summary, in the present article we have obtained some known results as that concerning to efficiency of metaphytes and other one we consider less known as it is the case for the negative values of the ecological function for photosynthesis, that is, with a dissipation term larger than the work produced as a possible explanation of the low overall efficiency of photosynthesis.

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Appendix

These data were taken or calculated from [11-13], [27-30]

1. Superior Plants and Cyanobacteria

$$\Delta G^{0} = 2880.31 \frac{\text{kJ}}{\text{mol}_{C_{6}H_{12}O_{6}}} \qquad \qquad \lambda = 680 \text{ nm}$$

2. Sulfur Purple Bacteria and Sulfur Green Bacteria, young bacteria

$$\Delta G^{0} = 429.64 \quad \frac{\text{kJ}}{\text{mol}_{C_{6}\text{H}_{12}\text{O}_{6}}} \qquad \lambda = 840 \text{ nm} \quad \lambda = 870 \text{ nm} \quad \lambda = 890 \text{ nm}$$

3. Sulfur Purple Bacteria, old bacteria

$$\Delta G^{0} = 744.57 \quad \frac{\text{kJ}}{\text{mol}_{C_{6}\text{H}_{12}\text{O}_{6}}} \qquad \qquad \lambda = 870 \text{ nm} \qquad \lambda = 890 \text{ nm}$$

4. Sulfur Purple Bacteria and Sulfur Green Bacteria

$$6\text{CO}_2 + 15\text{H}_2\text{O} + 3\text{Na}_2\text{S}_2\text{O}_3 \longrightarrow \text{C}_6\text{H}_{12}\text{O}_6 + 6\text{H}_2\text{O} + 6\text{Na}\text{HSO}_4$$
$$\Delta G^0 = 621.47 \frac{\text{kJ}}{\text{mol}_{\text{C}_6\text{H}_{12}\text{O}_6}} \qquad \lambda = 840 \text{ nm} \qquad \lambda = 870 \text{ nm} \qquad \lambda = 890 \text{ nm}$$

5. Non Sulfur Purple Bacteria and Non Sulfur Green Bacteria

$$6CO_2 + 12CH_3CH_2OH \longrightarrow C_6H_{12}O_6 + 12CH_3CHO + 6H_2O$$

$$\Delta G^0 = 584.86 \frac{kJ}{mol_{C_6H_{12}O_6}} \qquad \qquad \lambda = 840 \text{ nm} \qquad \lambda = 870 \text{ nm} \qquad \lambda = 960 \text{ nm}$$

6. Non sulfur Purple Bacteria and Non Sulfur Green Bacteria

$$\Delta G^{0} = 71.27 \quad \frac{\text{kJ}}{\text{mol}_{C_{6}\text{H}_{12}\text{O}_{6}}} \qquad \qquad \lambda = 840 \text{ nm} \quad \lambda = 870 \text{ nm} \quad \lambda = 960 \text{ nm}$$

7. Non Sulfur Purple Bacteria

$$6CO_2 + 12Succinic Ac. \longrightarrow C_6H_{12}O_6 + 6H_2O + 12Fumaric Ac.$$
$$\Delta G^0 = 1066.56 \frac{kJ}{mol_{C_6H_{12}O_6}} \qquad \qquad \lambda = 870 \text{ nm} \qquad \lambda = 960 \text{ nm}$$

8. Non Sulfur Purple Bacteria

$$6CO_2 + 12Malic Ac. \longrightarrow C_6H_{12}O_6 + 6H_2O + 12Oxalacetic Ac.$$
$$\Delta G^0 = 609.48 \frac{kJ}{mol_{C_6H_{12}O_6}} \qquad \lambda = 870 \text{ nm} \qquad \lambda = 960 \text{ nm}$$

9. Heliobacteria

$$3CH_{3}COOH + 6H_{2}O \longrightarrow 6CO_{2} + 12H_{2}$$
$$6CO_{2} + 12H_{2} \longrightarrow C_{6}H_{12}O_{6} + 6H_{2}O$$
$$\Delta G^{0} = 320.65 \frac{kJ}{mol_{C_{6}H_{12}O_{6}}} \qquad \lambda = 798 \text{ nm}$$

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