



## Metabolic model of *Arthrospira maxima* growth

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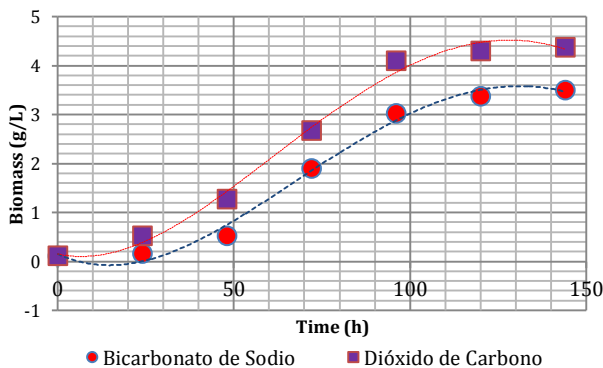
**Introduction.** The cyanobacterium *Arthrospira* has gained considerable attention worldwide as a source of several nutraceuticals (1). It has been commercially produced for more than 30 years under solar radiation in artificial ponds (2). An attractive alternative to reduce the carbon dioxide emissions could be the use of photosynthetic microorganisms able to utilize this pollutant as a carbon source for their growth. Thus, the photoautotrophic cultivation of *Arthrospira sp.* allows producing valuable biomass (3).

The objective is to design a metabolic model that can predict the growth rate and metabolites concentration at different growth conditions in a photobioreactor.

**Methods.** A summary of the most important reactions in the metabolism was made. The technique of extracting phycobiliproteins using modified phosphate regulator was used. The technique of extraction of chlorophyll with acetone 90% modified was used. Biomass was obtained by dry weight. Light is measured using a photodetector.

### Results.

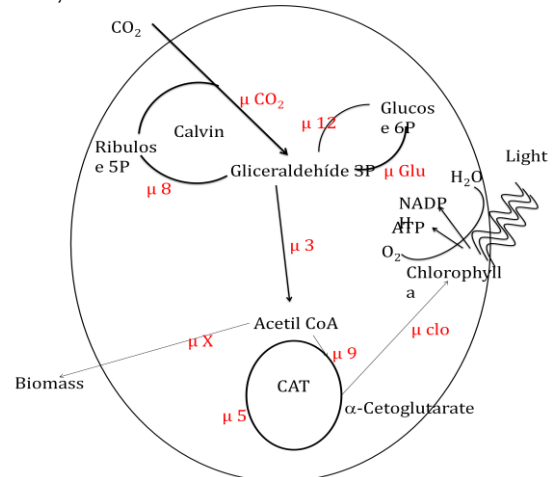
Araujo (4) used a novel design photobioreactor and found the intracellular concentration of phycocyanin and chlorophyll to increase in semi-continuous culture under constant agitation and irradiation with lamps that were not optimal for photosynthesis. Maya (5) developed a Venturi suction device for aspirating gas containing CO<sub>2</sub> for mixing and feeding, and found that the maximum specific growth rate for *Arthrospira maxima* ( $\mu_{max}$ ) to increase when using CO<sub>2</sub> as a carbon source compared with NaHCO<sub>3</sub>. Thus, the coupling system of the Venturi injector allows the photobioreactor to operate for optimal growth.



**Fig.1.** Biomass production rate rises significantly using carbon dioxide, compared with using sodium bicarbonate.

Figure 2 schematizes the photoautotrophic pathways network, lumping CO<sub>2</sub> uptake and Calvin cycle to produce Gly 3P, water photolysis to produce NADP and liberate O<sub>2</sub>, gluconeogenesis, and lower glycolysis to acetyl-CoA.

This metabolite is used for biomass and chlorophyll synthesis, ATP used for maintenance is also considered.



**Figure 2.** Outlining of the *Arthrospira maxima* metabolic model.

The stoichiometry of the lumped reactions is used to set up the conservation equations for Gly 3P, glucose, acetyl CoA, chlorophyll, NAD, ATP, as shown in the following equations:

$$\begin{aligned} \frac{d\epsilon ATP}{dt} &= -\frac{1}{3}\mu_{CO_2} + \mu_3 - \frac{9}{55}\mu_4 - \frac{1}{3}\mu_{12} - \frac{1}{5}\mu_8 - Y\mu_6 - m_{ATP} + \phi_{ATP} = 0 \\ \frac{d\epsilon Ru5P}{dt} &= -\frac{5}{6}\mu_{CO_2} + \mu_8 = 0 \\ \frac{d\epsilon Gly\ 3P}{dt} &= -\frac{3}{2}\mu_3 - \mu_8 - \mu_{Glu} + \mu_{CO_2} + \mu_{12} = 0 \\ \frac{d\epsilon CO_2}{dt} &= -\frac{1}{6}\mu_{CO_2} + \frac{1}{2}\mu_3 + \frac{1}{5}\mu_9 - \frac{1}{4}\mu_5 + \frac{6}{55}\mu_{Clor} - \frac{1}{10}\mu_x = 0 \\ \frac{d\epsilon Oxalacetato}{dt} &= \mu_5 - \frac{4}{5}\mu_9 = 0 \\ \frac{d\epsilon Glu}{dt} &= \mu_{glu} - \frac{1}{2}\mu_{12} \\ \frac{d\epsilon NAD}{dt} &= -\mu_3 - \frac{1}{5}\mu_9 - \frac{1}{2}\mu_5 + \frac{8}{55}\mu_{Clor} - \frac{1}{10}\mu_x + \phi_{NAD} = 0 \\ \frac{d\epsilon Ac\ CoA}{dt} &= \mu_3 - \frac{2}{5}\mu_9 - \mu_6 = 0 \end{aligned}$$

**Conclusions.** It was demonstrated that the cellular growth is better using CO<sub>2</sub> as carbon source than using sodium bicarbonate.

A novel experimental set up for photobioreactor operation is described.

### References

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