INFLUENCE OF ENVIRONMENTAL CONDITIONS ON CHRYSOCHROMULINA ROTALIS FOR THE PRODUCTION OF BIOACTIVE COMPOUNDS AND HIGH-VALUE COMPOUNDS

IX SIMPOSIO de Investigación **EN CIENCIAS EXPERIMENTALES**

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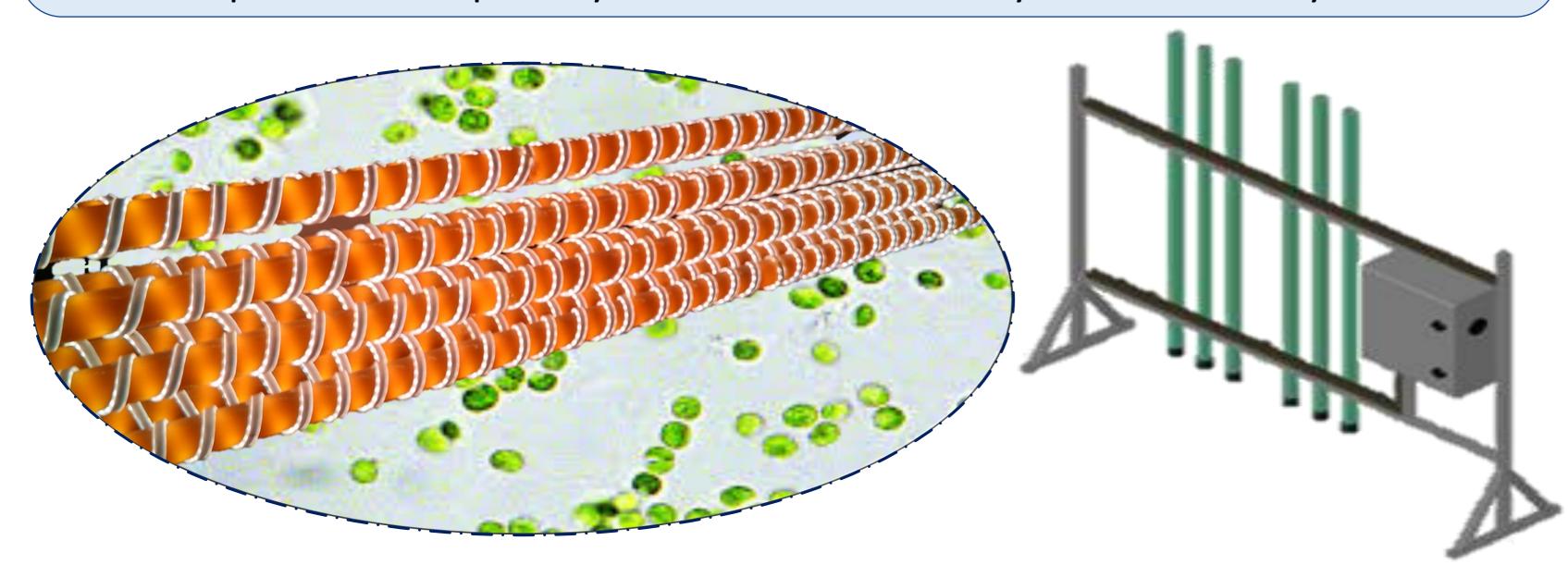
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INTRODUCTION

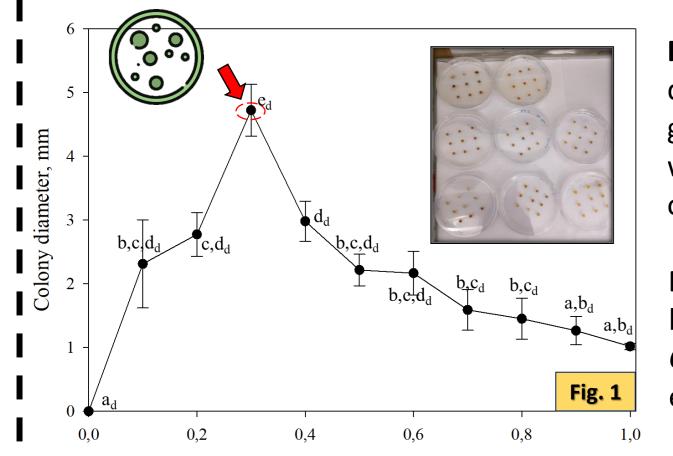
Identification of appropriate environmental culture conditions is a prerequisite for the maximization of microalgae biomass and bioactive production. Phylum haptophyta includes some flagellated microalgae known to be producers of high value biomolecules. Chrysochromulina rotalis, a type of haptophyte, has previously been cultivated on a small scale and has been shown to be a producer of carotenoids as fucoxanthin^[1], fatty acids as DHA^[2] and bioactive compounds^[3]. However, a comprehensive optimization process of culture conditions has not been performed yet. In addition, most flagellated microalgae are difficult to maintain for a long time in cell banks since they usually do not allow cultivation in a solid medium. This, also makes it difficult to obtain monoalgal cultures.

OBJECTIVES

The target of this work is to optimize the operational environmental conditions (salinity, irradiance and temperature) for the cultivation of *C. rotalis*, which maximize the production of high added value products and bioactive compounds, cultivated photoautotrophically and in an economically sustainable way.



RESULTS AND DISCUSSION



Agar percentage, % v/v

Figure 1. Growth in diameter of *C. rotalis* grown in Petri dishes (\$\frac{\frac{1}{2}}{2}\$) with variable proportions of technical agar.

Figure 2. Results of the biomass productivity of C. rotalis in mg/L·day for each salinity value.

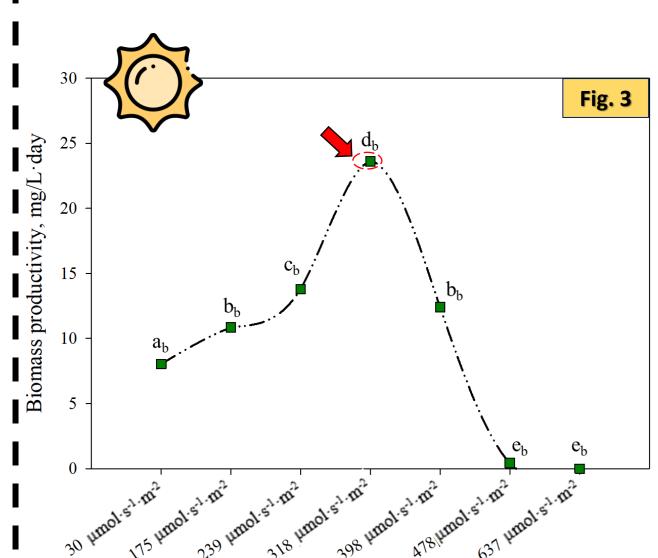
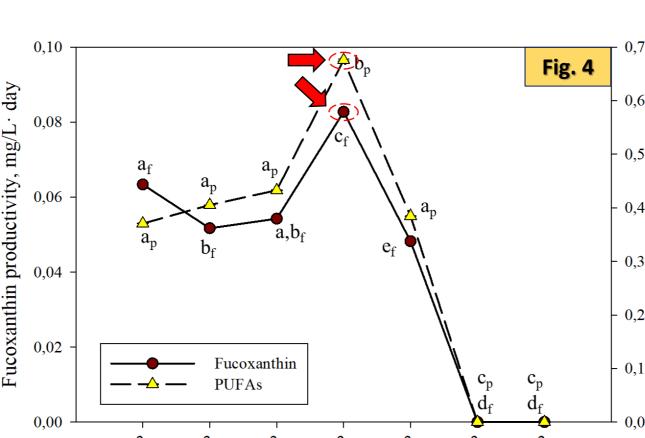


Figure Biomass productivity of *C. rotalis* ਵੇਂ in mg/L·day for each \frac{1}{20} irradiance value.

Figure 4. Productivity of **§** fucoxanthin and PUFAs of C. rotalis in mg/L·day for each irradiance value.



MATERIAL AND METHODS

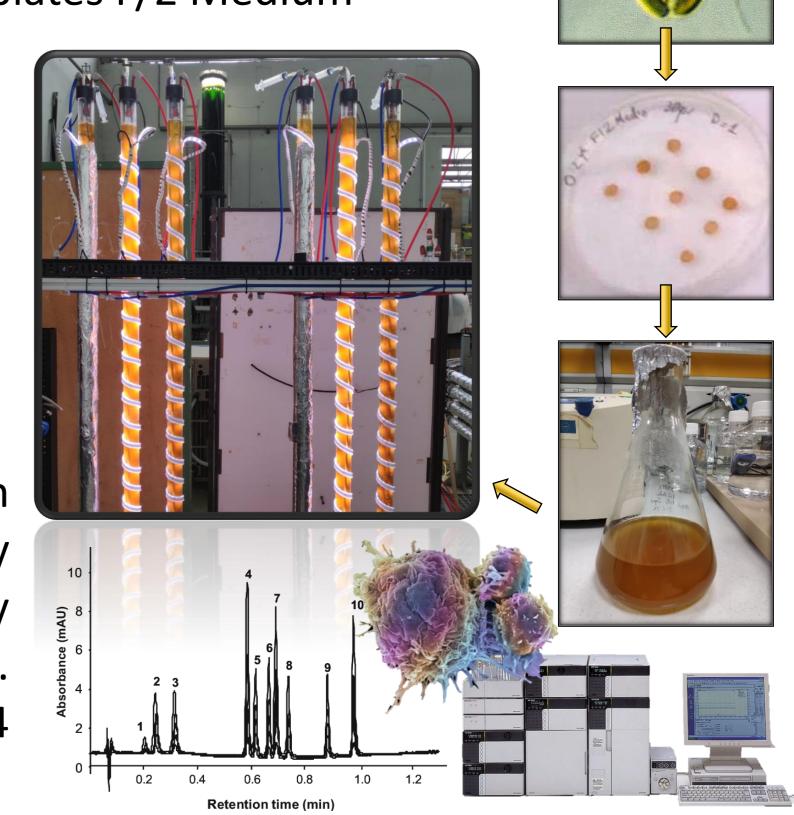
Biomass: Chrysochromulina rotalis (UPV)

Strain maintenance: Petri plates F/2 Medium Agar $(0,1-1)\%^{[4]}$.

Photobiorreactor:

2,5L Bubble column with LEDs^[5] (30-637 μ mol·m⁻²·s⁻¹), pH 8,5 at (15–35)±1°C in discontinuous mode.

Biomass characterization: Carotenoids by HPLC-DAD (Cerón García et al., (2018)^[6] and fatty gas chromatography (Rodríguez Ruiz et al., (1998)^[7]. Antiproliferative activity against 4 tumor lines.



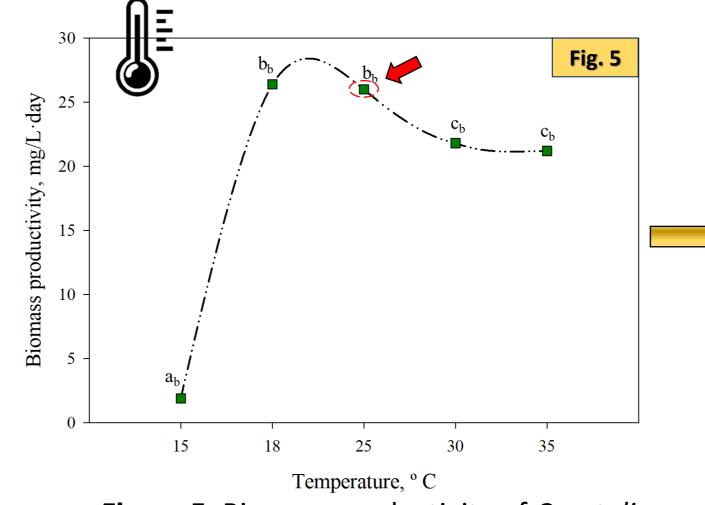


Figure 5. Biomass productivity of *C. rotalis* in mg/L·day for each temperature value.

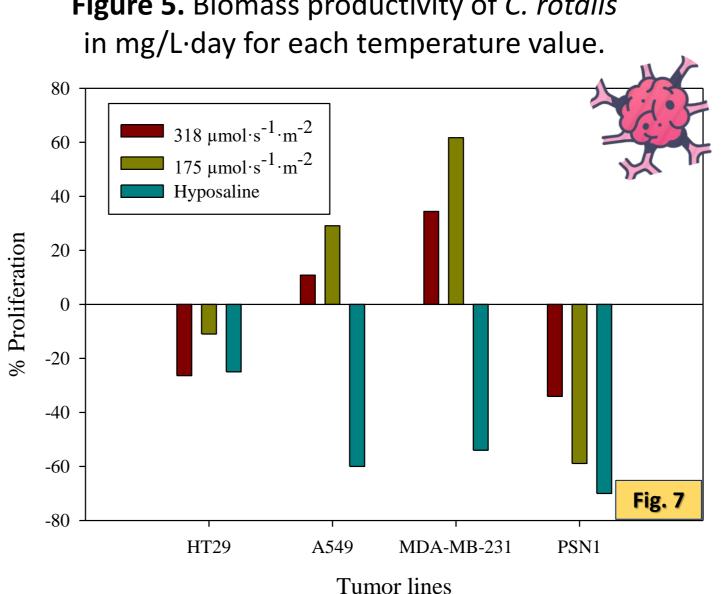


Fig. 6 — Fucoxanthin Temperature, °C

Figure 6. Productivity of fucoxanthin and PUFAs of C. rotalis in mg/L·day for each temperature value.

Figure 7. Percentage of proliferation of 4 types of human tumor cells in assays using methanolic extracts of biomass obtained in the optimization of average irradiance and hyposaline shock. A negative percentage indicates a recess in proliferation due to cytotoxicity.

HT-29 (colon adenocarcinoma) A549 (lung carcinoma) PSN-1 (pancreatic adenocarcinoma) MDA-MB-231 (breast adenocarcinoma)

CONCLUSIONS

- Optimal growth value for cultivation and maintenance in Petri dishes for the microalgae C. rotalis in the 0.3% agar condition, not described to date for any species of haptophyte.
- ✓ Optimal productivity at 30PSU, 318 μmol·s⁻¹·m⁻² of I_{avg} using LEDs and 25°C temperature \implies P_{biomass}: 13, 23.6, 26 mg/L·day, respectively.
- ✓ Finally, a **high antitumor and antiproliferative capacity** has been demonstrated due to the cytotoxicity of the tested biomasses, obtained under hyposalinity and optimal irradiance conditions.
- ✓ Thus, *C. rotalis* is proposed as a haptophytic microalgae that could be scaled at an industrial level, with significant applications in the pharmaceutical industry and the fight against human tumor cells.

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