Supercritical carbon dioxide extraction of biological compounds from microalgae

Beatriz P. Nobre¹, Luisa Gouveia¹, Luca Campenni¹², Filipa M. Marcelo³, António F. Palavra², Rui L. Mendes¹

1- Unidade de Bioenergia, LNEG, Estrada do Paço do Lumiar 22, 1649-038 Lisboa, Portugal. Phone: +351217127213, Fax: +351217163636, e-mail: Beatriz.nobre@mail.ineti.pt, Luisa.gouveia@ineti.pt, rui.mendes@ineti.pt
2- IST, Centro Química Estrutural, DEQB, Av Rovisco Pais 1049-001, Lisboa, Portugal, Phone. +351218419387, Fax: +351218464455, e-mail: Antonio.palavra@ist.utl.pt
3- Centro de Investigaciones Biológicas, CIB, c/Ramiro de Maeztu 9, 28040 Madrid, Spain, Phone: +34 918373112, e-mail: fmmarcelo@cib.csis.es

Microalgae are eukaryotic, photosynthetic, unicellular microorganisms that present a great genetic diversity. These microorganisms produce several biological compounds that are important in the nutraceutical and pharmaceutical industries, such as hydrocarbons, carotenoids, vitamins, polyunsaturated fatty acids, etc [1]. Supercritical fluid extraction (SFE) of biological compounds from microalgae has some advantages over the conventional solvent extraction methods, because the compounds can be obtained without contamination by the organic solvent and thermal degradation [2]. On the other hand, it is possible a high efficiency of the extraction and the selectivity for certain compounds is more easily achieved with SFE than with organic solvent extraction. Moreover, supercritical carbon dioxide extraction has also the advantage of using a non-toxic, non-flammable and cheap solvent.

Supercritical fluid extraction studies of biological compounds from several microalgae, such as Botryococcus braunii, Chlorella vulgaris, Dunaliella salina, Haematococcus pluvialis and Arthospira (Spirulina) maxima have been carried out in our laboratories [3,4,5,6,7]. A semi-continuous laboratorial-scale supercritical fluid extraction apparatus, which allows working in the temperatures range of 40-80 ºC and pressures up to 400 bar, was used to perform all studies [8]. The extracted compounds were identified and quantified off-line by HPLC, GC and UV/Visible spectrophotometry. The effect of temperature, pressure and solvent flow-rate, together with the use of co-solvent and pre-treatment of the microalgae biomass, was assessed in these extraction studies.

The aim of this work is to present an overview of the most important results achieved during the supercritical carbon dioxide extraction of biological compounds from the several microalgae studied with special emphasis on the effect of the pre-treatment of the biomass and the use of entrainers in the extraction of carotenoids.

References