



DITERPENES OF BROWN ALGAE *Canistrocarpus cervicornis* FROM CEARÁ

Ana Cláudia Philippus¹, Lucas Felipe de Oliveira Vieira¹, Gabriele Andressa Zatteli¹, Efstathia Ioannou², Vassilios Roussis², Miriam de Barcellos Farkenber¹

¹Universidade Federal de Santa Catarina, Florianópolis/SC, Brasil; ²University of Athens, Athens, Greece;
e-mail address: anaphilippus@gmail.com

Abstract: *Canistrocarpus cervicornis* is widely distributed in the world [1]. Previous investigations led to isolation of dolastane and secodolastane diterpenes, for which there are many reported biological activities [2,3,4,5]. Marine organisms present usually a considerable chemical diversity among samples from different locations [6]. Therefore, since samples from “Pedra do Paraíso” (CE/Brazil) were not previously investigated, the aim of this project was to isolate secondary metabolites from this selected sample. Algal material was dried under cold air and soaked with dichloromethane/methanol 2:1. Fractionation of the crude extract was performed by successive chromatographic separations with silica gel and gradient of solvents with increasing polarity. Vacuum Liquid Chromatography (VLC) was used in the first step; selected fractions were submitted to column chromatography with medium pressure and, eventually, to High Performance Liquid Chromatography (HPLC). Samples considered pure by Thin Layer Chromatography (TLC) analysis, were analyzed by Nuclear Magnetic Resonance (NMR) in deuterated chloroform in spectrometers Bruker AC200 and Bruker DRX400. Two pure compounds were obtained (CC1 and CC2). The ¹H NMR spectrum of CC1 includes four singlets between 0.83 and 1.24 ppm, for hydrogens of the methyl groups; signals for a deshielded hydrogen at 5.52 ppm, corresponding to one olefinic H; and at 4.90 and 4.80 ppm, corresponding to the exomethylene H atoms. CC2 presented a similar spectrum, but with signals for two olefinic hydrogens. The database Marinlit® allowed the identification of these compounds as (4*S*,9*R*,14*S*)-4-acetoxy-9,14-dihydroydolasta-1(15),7-diene (CC1) and (4*S*,14*S*)-4-acetoxy-14-hydroxydolasta-1(15),7,9-triene (CC2), tricyclic diterpenes of the dolastane type, confirming the relative abundance of these diterpenes among Dictyotaceae. CC1 presents several biological activities, especially antiviral [4], antileishmania [5], anticoagulant, antiplatelet [2], among others. For CC2, no biological activity was reported so far. Additional studies aiming isolation of other metabolites are in progress.

References:

- [1] Guiry, M. D.; Guiry, G. M. 2015. AlgaeBase. World-wide electronic publication, National University of Ireland. Available in: <<http://www.algaebase.org>>. Access: 11 jun 2015.
 - [2] Moura, L. A., Bianco, E.M., Pereira, R.C., Teixeira, V.L., Fuly, A.L. 2011. Anticoagulation and antiplatelet effects of a dolastane diterpene isolated from the marine brown alga *Canistrocarpus*. *J. Thromb. Thrombolysis*. 31: 235–240.
 - [3] Teixeira, V. L.; Tomassini, T., Fleury, B.G., Kelecom, A. 1986. Dolastane and secodolastane diterpenes from the marine brown alga *Dictyota cervicornis*. *J. Nat. Prod.* 49: 570–575.
 - [4] Vallim, M. A., Barbosa, J.E., Cavalcanti, D.N., De-Paula, J.C., Silva, V.A.G.G., Teixeira, V.L., Paixão, I.C.N.P. 2010. *In vitro* antiviral activity of diterpenes isolated from the brazilian brown alga *Canistrocarpus cervicornis*. *J. Med. Plants Res.* 4: 2379-2382.
 - [5] Santos, A.O., Britta, E. A., Bianco, E.M., Ueda-Nakamura, T., Dias Filho, B.P., Pereira, R.C., Nakamura, C.V. 2011. 4-Acetoxydolastane diterpene from the brazilian brown alga *Canistrocarpus cervicornis* as antileishmanial agent. *Mar. Drugs*. 9: 2369-2383.
 - [6] Stengel, D. B., Connan, S., Popper, Z.A. 2011. Algal chemodiversity and bioactivity: sources of natural variability and implications for commercial application. *Biotechnol. Adv.* 29: 483-501.
-