CARBONERO, E. R. ; <u>RUTHES, A. C.</u>; WAIHRICH, L. G. ; AMAZONAS, M. A. L. A. ; <u>GORIN, P. A. J.</u>; <u>IACOMINI, M.</u>. Further investigations on the polysaccharides obtained from the stems and caps of Macrocybe titans. In: XXXV Reunião Anual da Sociedade Brasileira de Bioquímica e Biologia Molecular (XXXV SBBq), 2006, Águas de Lindóia. Programa e Resumos da XXXV Reunião Anual, 2006.

http://sbbq.iq.usp.br/arquivos/2006/cdlivro/resumos/R8199.html

XXXV Reunião Anual da SBBq

ResumoID:8199

## Further Investigations on the Polysaccharides Obtained from the Stems and Caps of Macrocybe titans

Carbonero, E. R.<sup>a</sup>; Ruthes, A. C.<sup>a</sup>; <u>Waihrich, L. G.</u><sup>a</sup>; Amazonas, M. A. L. A.<sup>b</sup>; Gorin, P. A. J.<sup>a</sup>; Iacomini, M.<sup>a</sup>

<sup>a</sup> Departamento de Bioquímica e Biologia Molecular, Universidade Federal do Paraná, UFPR,
C.P. 19046, CEP 81531-990, Curitiba, PR, Brasil; <sup>b</sup> Centro Nacional de Pesquisa de Florestas,
EMBRAPA-Florestas, Caixa Postal 319, CEP 83411-000, Colombo, PR, Brasil

e-mail: lariwaihrich@hotmail.com

The giant mushroom Macrocybe titans have drawn the attention on account of their food potential. A chemical analysis was carried out on its stems and caps, in order to determine their carbohydrate structures. Each portion was extracted with CHCl<sub>3</sub>-MeOH and then MeOH-H<sub>2</sub>O to remove low-molecular weight compounds. Each resulting residue was submitted, successively, to aqueous extraction (x 6) and 2% aq. KOH (x 6) at 100°C, and the extracted polysaccharides, which were recovered by ethanol precipitation (WP, KP, WS, and KS fractions), were dialyzed against tap water for 24 h, and freeze-dried. The polysaccharide fractions were then submitted to purification procedures by freeze-thawing to give insoluble (I- WP, I-KP, I-WS, and I-KS fractions) and water-soluble fractions (S-WP, S- WS, S- KP, and S-KS fractions). According to the GC-MS analysis of derived alditol acetates, all the insoluble-water fractions had glucose as its main monosaccharides, while those of soluble fractions were Gal, Glc, Fuc, and Man. Due to their similarity, as shown by GC-MS and <sup>13</sup>C-NMR, the soluble fractions obtained from stems (S-WS, S-KS) and caps (S-WP, S-KP) these were combined, furnishing fractions SW and SK. SW and SK fractions were treated with Fehling solution, giving supernatants (FS-SW and FS-SK) and precipitates (FP-SW and FP-SK). FS-SW, FS-SK and FP-SK fractions had Glc as their main monosaccharide while FP-SW contained Fuc, Man, Gal and Glc. The <sup>13</sup>C-NMR spectrum of the FP-SK fraction showed six main signals ( $\delta$  103.2, 76.1, 75.3, 73.4, 70.2, and 69.2, corresponding to C-1, C-3, C-5, C-2, C-4, and C-6) of a  $\beta$ -(1 $\rightarrow$ 6)-linked glucan. Fractions FP-SW, FS-SW, and FS-SK gave rise to heterogeneous (HPSEC), so that they are been purified by ultrafiltration membranes (300, 30 and 10 kDa Mr cut-offs). Additional structural studies of these fractions are being carried out by the above methods.

Supported by CAPES, CNPq, PRONEX- Fundação Araucária