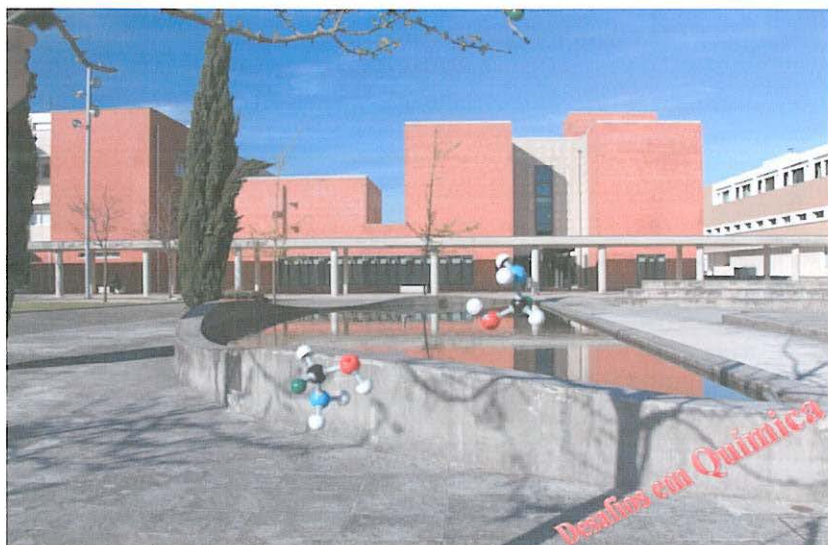


XXIII Encontro Nacional da SPQ



Aveiro 12 a 14 de Junho de 2013



Titulo

*Livro de Resumos do XXIII Encontro Nacional da SPQ
Desafios em Química*

Coordenadores

Diana C. G. A. Pinto e Artur M. S. Silva

Fotografias

José M. G. Pereira

Fotografia da Capa

Departamento de Química e Laboratórios Tecnológicos da Universidade de Aveiro

Edição

*Sociedade Portuguesa de Química
Av. da República, 45 - 3º Esq.
1050-187 Lisboa - Portugal*

Data

Junho de 2013

Tiragem

350 Exemplares

Depósito Legal

360265/13

Impressão e acabamentos

Sersilito-Empresa Gráfica, Lda.

Catálogo recomendada

*Livro de resumos do XXIII Encontro Nacional da SPQ - Desafios em Química
Universidade de Aveiro, 2013 – 387 p.*

Química - Congressos

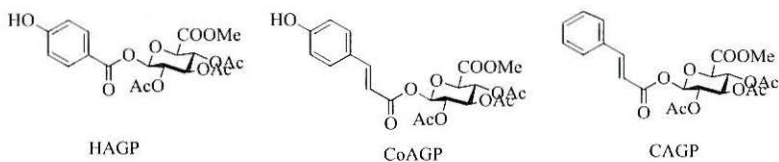
Reservados todos os direitos. Esta publicação não pode ser reproduzida ou transmitida, no todo ou em parte, por qualquer processo, eletrónico, mecânico, fotocópia, gravação ou outros, sem prévia autorização do Editor.

Growth inhibitory activity of *Coprinopsis atramentaria* extract, phenolic acids, cinnamic acid and protected acetylated glucuronides on human tumor and non-tumor cell lines

Sandrina A. Heleno,^{a,b} Isabel C.F.R. Ferreira,^b Ricardo C. Calhella,^{a,b} Anabela Martins,^b Ana P. Esteves,^a Maria João R.P. Queiroz^a

^aCentro de Química, Universidade do Minho, Campus de Gualtar 4710-057 Braga, Portugal; ^bCentro de Investigação de Montanha (CIMO), ESA, Instituto Politécnico de Bragança, Campus de Santa Apolónia, apartado 1172, 5301-855 Bragança, Portugal.

Mushrooms represent an important source of compounds with antitumor and immunostimulating properties.^[1] Our attention is being focused on the capacity of phenolic acids from mushrooms to inhibit the growth of different tumor cell lines.^[2] *Coprinopsis atramentaria* (Bull.: Fr.) Redhead, Vilgalys & Moncalvo, is a wild edible mushroom previously characterized by us for its nutritional composition, and also showed a notable antioxidant activity; *p*-hydroxybenzoic (4.71 mg/100 g dry weight), *p*-coumaric (0.82 mg/100 g) and cinnamic (1.70 mg/100 g) acids were identified in its phenolic extract.^[3] Herein, the growth inhibitory activity on human tumor cell lines (breast- MCF-7, non-small cell lung- NCI-H460, colon- HCT-15, hepatocellular- HepG2 and cervical- HeLa) and non-tumor porcine liver primary cell culture (PLP2), was evaluated by sulforhodamine B assay, using *C. atramentaria* extract, phenolic acids, cinnamic acid and their protected glucuronides (HAGP, CoAGP and CAGP; Scheme) obtained reacting *p*-hydroxybenzoic (HA), *p*-coumaric (CoA) and cinnamic (CA) acids, respectively, with acetobromo- α -D-glucuronic acid methyl ester.^[4]



C. atramentaria extract proved to have high antitumor potential against NCI-H460 ($GI_{50}=15.13 \pm 1.35 \mu\text{g/mL}$), HCT-15 ($GI_{50}=36.44 \pm 3.30 \mu\text{g/mL}$) and MCF-7 ($GI_{50}=53.10 \pm 4.72 \mu\text{g/mL}$) cell lines, without toxicity for non-tumor porcine liver primary cells ($GI_{50}>400 \mu\text{g/mL}$). The protected compounds HAGP, CoAGP and CAGP revealed higher antitumor activity when compared with the parental acids for all the tested cell lines. The most active compound was HAGP against HepG2 cell line ($GI_{50}=57.78 \pm 5.65 \mu\text{M}$).

Acknowledgments: FCT for financial support to the Portuguese NMR network and COMPETE/QREN/EU (PTDC/AGR-ALI/110062/2009, PEst-OE/AGR/UI0690/2011 and PEst-C/QUI/UI0686/2011) for financial support. S.A. Heleno (BD/70304/2010) and R.C. Calhella (BPD/68344/2010) also thank FCT, POPH-QREN and FSE.

References

- [1] Ferreira, I.C.F.R.; Vaz, J.A.; Vasconcelos, M.H.; Martins, A. *Anticanc. Ag. Med. Chem.* **2010**, *10*, 424.
- [2] Vaz, J.A.; Almeida, G.M.; Ferreira, I.C.F.R.; Martins, A.; Vasconcelos, M.H. *Food Chem.* **2012**, *132*, 482.
- [3] Heleno, S.A.; Barros, L.; Queiroz, M.-J.R.P.; Santos-Buelga, C.; Ferreira, I.C.F.R. *J. Agric. Food Chem.* **2012**, *60*, 4634.
- [4] Heleno, S. A.; Ferreira, I. C. F. R.; Esteves, A. P.; Ciric, A.; Glamoclija, J.; Martins, A.; Sokovic, M.; Queiroz, M.-J. R. P. *Food Chem. Toxicol.* **2013**, *58*, 95.