



Abstract book

The eighth international workshop
on edible mycorrhizal mushrooms

IWEMM8



The sixth conference of the *Tuber aestivum/uncinatum*
European scientific group

TAUESG 6

Livre des résumés
Congrès international
sur les champignons mycorrhiziens comestibles

October 10th - 17th 2016

10-17 octobre 2016

Espace de congrès Clément-Marot

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Topic of the congress (Les thèmes du congrès)

1. Culture
2. Biologie
3. Ecologie
4. Taxonomie
5. Biologie moléculaire
6. Changement climatique
7. Développement de l'économie rurale
8. Mycotourisme, agritourisme et gastronomie
9. Chimie alimentaire et bénéfices sur la santé
10. Conservation et protection de l'environnement
11. Gestion des populations en milieu naturel et cultivé
12. Autres questions relatives aux champignons mycorrhiziens comestibles

Scientific Committee (Le Comité scientifique international)

- Shannon Berch - Canada
- Carolina Barroetaveña - Argentine
- Dominique Barry - France
- Gérard Chevalier - France
- Simon Egli – Suisse
- Arzu Roberto Flores - Guatemala
- Alexis Guerin-Laguette – Nouvelle-Zélande, Secretary general
- Ian Hall – Nouvelle-Zélande
- Lahsen Khabar – Maroc
- Fernando Martínez-Peña - Espagne
- Jesus Pérez-Moreno - Mexique
- Asun Morte - Espagne
- Daniel Mousain - France
- Claude Murat - France
- David Pilz - USA
- Jean-Marc Olivier - France
- Jean Rondet - France
- Marc-André Selosse - France
- Pierre Sourzat - France,
- Aziz Türkoğlu - Turquie
- Wang Yun - Nouvelle-Zélande, Chine
- Akiyoshi Yamada - Japon
- Yu Fuqiang - Chine
- Alessandra Zambonelli - Italie



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CONTRIBUTION OF PHENOLIC ACIDS TO THE ANTI-INFLAMMATORY ACTIVITY OF EDIBLE MYCORRHIZAL MUSHROOMS

Oludemi Taofiq^{ab}, Ricardo C. Calhella^{ac}, Sandrina A. Heleno^{ac}, Lillian Barros^a, Anabela Martins^a, Celestino Santos-Buelga^b, Maria João R.P. Queiroz^c, Isabel C.F.R. Ferreira^a
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Edible mycorrhizal mushrooms are rich in several bioactive compounds responsible for their interesting health benefits (1). Polysaccharides, terpenoids and phenolic compounds have been indicated as the most important contributors to the anti-inflammatory activity of mushroom species (2). In the present study, ethanolic extracts of seven mycorrhizal mushrooms were obtained by maceration and chemically characterized in terms of phenolic acids by HPLC-DAD. The extracts, the identified phenolic acids and also synthesised methylated and glucuronated derivatives of those acids were studied regarding anti-inflammatory activity evaluated through inhibition of nitric oxide (NO) in LPS (lipopolysaccharide) activated RAW 264.7 macrophages. Among the studied mushrooms, *Boletus impolitus* presented the highest amount of phenolic acids ($675 \pm 23 \mu\text{g/g}$), due to the high contribution of cinnamic acid ($505 \pm 12 \mu\text{g/g}$), and also displayed the highest NO production inhibition ($\text{EC}_{50} = 166 \pm 10 \mu\text{g/mL}$). The phenolic acids content in the different species was as follows: *B. impolitus* > *C. cibarius* > *A. caesaria* > *L. deliciosus* > *B. aereus* > *M. esculenta* > *B. edulis*, while the NO production inhibition EC_{50} values were in the order: *B. impolitus* > *A. caesaria* > *C. cibarius* > *L. deliciosus* > *M. esculenta* > *B. aereus* > *B. edulis*. Among the individual compounds, cinnamic acid (CA) showed the highest activity ($\text{EC}_{50} = 182 \pm 16 \mu\text{M}$), followed by *p*-hydroxybenzoic acid (HA; $239 \pm 29 \mu\text{M}$) and *p*-coumaric acid (CoA; $442 \pm 33 \mu\text{M}$), which highlights the contribution of cinnamic acid for the anti-inflammatory potential displayed by mushrooms. Comparing the NO production inhibition activity of the parent phenolic acids and their conjugated derivatives (G-glucuronated, M-methylated), the order was as follows: *p*-hydroxybenzoic acid: HA > HA-M3 > HA-M2 > HA-M1 > HA-G; *p*-coumaric acid: CoA-M1 > CoA-G > CoA-M2 > CoA-M3 > CoA and cinnamic acid: CA-G > CA > CA-M. The results in the present study show the contribution of phenolic acids to the anti-inflammatory activity of mushrooms and also demonstrated that conjugation reactions can influence the bioactivity of the phenolic acids.

Reference

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