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A087 Composition and antioxidant activity in vitro of the essential oil of *Thymus bracteosus* in comparison to *Thymus vulgaris**Ozren Ercegova*^a, *Hartwig W. Pfeifhofer*^b, *Zeljka Males*^a, *Misko Plazibat*^c and *Adelheid H Brantner*^d^a Department of Pharmaceutical Botany, Faculty of Pharmacy and Biochemistry, University of Zagreb, Schrottova 39, 10000 Zagreb, Croatia. ^b Institute of Plant Physiology, University of Graz, Schubertstrasse 51, A-8010 Graz, Austria. ^c Department of Botany, Faculty of Science, University of Zagreb, Maculicev trg 20/II, 10000 Zagreb, Croatia; ^d Institute of Pharmacognosy, University of Graz, Universitaetsplatz 4/1, A-8010 Graz, Austria.

The composition of the essential oil and the antioxidant properties of the dried herbs of *Thymus bracteosus* Vis. ex Benth (Lamiaceae), a plant endemic to the Dinaric Karst, were investigated to find out if *T. bracteosus* can be used as herbal drug in the same way as *T. vulgaris*. The composition of the essential oil was analysed by GC/FID and GC/MS techniques. A data base was used for automatic identification of GC/MS peaks, linear retention indices were compared with published data or authentic compounds. GC/MS analysis of *T. bracteosus* essential oil (yield after hydrodistillation 0.06%) revealed the presence of 83 compounds. Out of them 65 substances were identified, representing 92% of the total components. The major compounds of the oil are α -pinene (6.3%), myrcene (7.1%), β -caryophyllene (9.6%), trans- β -farnesene (6%) and germacrene-D (11.4%). Analysis of *T. vulgaris* oil (yield after hydrodistillation 2.5%) showed the presence of 80 compounds, 64 of which were identified, representing 96.8% of the total compounds. The main constituents are *p*-cymene (37.8%) and thymol (36.7%). Differences of the essential oils of the two investigated plant species were also caused by the variation of the content of monoterpenes (*T. bracteosus* 28.9%, *T. vulgaris* 95.1%) and sesquiterpenes (*T. bracteosus* 62.2%, *T. vulgaris* 1.1%). As *T. bracteosus* is not fulfilling the requirements of the pharmacopoeias (DAB, OEAB, Ph. Helv.: yield at least 1.2% and 1.5% resp.) it cannot be used pharmaceutically in the same way as *T. vulgaris*. The essential oils of the two *Thymus* species were also investigated on their radical scavenger capacity measuring photometrically the disappearance of DPPH* (1) (*T. vulgaris* IC₅₀ 15.88 μ g/ml, reference rutin IC₅₀ 3.01 μ g/ml). Furthermore, the ability of the test samples to inhibit peroxidation of membrane lipids was tested (2) (*T. vulgaris* IC₅₀ 44.20 μ g/ml, reference fisetin IC₅₀ 7 μ g/ml). The essential oil of *T. bracteosus* did not show any antioxidant activity in both test systems.

References: 1. Hatano T. et al. (1988) Chem. Pharm. Bull. 36: 2090-2097. 2. Houghton P.J. et al. (1995) Planta Med. 61: 33-36.**A088 Lignicolous fungi as potential natural antioxidants***N. Mimica-Dukić*^a, *M. Karaman*^b, *M. Matavulj*^b, *R. Pavkov*^c and *M. Popović*^a^a Institute of Chemistry, Faculty of Natural Sciences, University of Novi Sad, 21 000 Novi Sad, Trg Dositeja Obradovića 3. Yugoslavia. ^b Institute of Biology, Faculty of Natural Sciences, University of Novi Sad, 21 000 Novi Sad, Trg Dositeja Obradovića 5. Yugoslavia. ^c Pharmaceutical Co. HEMOFARM, 13 000 Vršac, Yugoslavia.

In the last decade higher (Basidiomycetes) fungi became of great importance as sources of pharmacological active substances. Among them, lignicolous fungi are found to be of particular medicinal significance. Although a wide variety of biological activities of fungi were evaluated (1), their antioxidant ability have not been examined so far. In the present study, the effect of following lignicolous fungi: *Ganoderma applanatum*, *Ganoderma lucidum*, *Meripilus giganteus* and *Flammulina velutipes* on the Fe²⁺/ascorbate induced lipid peroxidation (LP) and free radical production is investigated. In the experiment MeOH and CHCl₃ extracts of dry fungi scorocarpus were used. MeOH extract of *G. applanatum* (10 mg/ml) exhibited highest inhibitory effect (61.52%) on LP in liposomes. All CHCl₃ extracts were less potent in reducing LP. The inhibitory activity was in dose dependent manner. Free radical scavenging capacity (RCS) was evaluated by following the effect of fungi extracts on OH radicals, generating in Fenton reaction (2), and measuring their ability to neutralize DPPH (2,2-diphenyl-1-picrylhydrazil) stable radical form, and transform it into reduced form (3). Although the RSC on OH radicals was very low, examined extracts exhibited very high DPPH scavenging activity. Highest DPPH-RSC was obtained with MeOH extract of *G. lucidum* (IC₅₀= 7.5 μ g/ml) and *G. applanatum* (IC₅₀= 10.3 μ g/ml), the lowest activity was obtained with *F. velutipes* (IC₅₀= 300 μ g/ml). CHCl₃ extracts of both *Ganoderma* also expressed strong DPPH-RSC (IC₅₀= 16.25 μ g/ml and 19.00 μ g/ml, respectively).

References: 1. Wasser, S.P. and Weis, A.L. (1999) Intern. J. Med. Mushrooms, 1: 31-62. 2. Cheesman, K.H. et al. (1988) Biochem. J. 252: 649-653. 3. Soler-Rivas, C. et al. (2000) Phytochem. Anal. 11: 330-338.