The Study on the Possibility of Using Ecological Materials with Antifungal Properties for Treating Natural Leathers

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Synthetic fungicides presently used are potentially harmful to both human health and for the environment. Recent research aims to fully or partially replace potentially toxic biocides with environmentally friendly materials. Essential oils extracted from plants may be an alternative to conventional fungicides. The purpose of the study was to investigate the possibility of using essential oil of cloves as alternative preservatives for skin treatment. Essential oil isolated from cloves (Eugenia caryophyllata) containing: eugenol – 78.03%, aceteugenol – 10.93%, caryophylene – 9.46%. Following the study, it is concluded that the selected essential oil of cloves can be used as an antifungal agent in the field of natural leather processing.

Keywords: natural leather, gas chromatography mass spectrometry (GC/MS), FT-IR spectrometry, UV-VIS spectrometry, Aspergillus niger

Experimental part

Materials and methods
- Cloves essential oil, Eugenia caryophyllata, (Adams, Romania) containing: eugenol – 78.03%, aceteugenol – 10.93%, caryophylene – 9.46% and alpha humulene – 1.08% etc.
- Ethanol (Chemical Company, Germany), colorless liquid, boiling point 78.37°C, density 0.79 g/cm³;
- The Box bovine leathers natural grain assortments, mineral tanned and wet finished by retanning, fatliquoring and dyeing (1.2-1.4 mm thick, dyed brown) (National Research and Development Institute for Textiles and Leather – Division Leather and Footwear Research Institute Bucharest, Romania).
- Biologic Material: Aspergillus niger.

Gas Chromatography Mass Spectrometry (GC/MS) Analysis:

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Analysis of the essential oils carried out by using Agilent 7890 A GC System equipped with Agilent 5795 C MS, and HP-5 MS (0.25 mm x 30 m i.d., film thickness 0.25). The carried gas helium (99.9%) at a flow rate of 1 mL/ min; ionization energy was 70 eV. Mass range m/z 50-650 amu. Data acquisition was scan mode. MS transfer line temperature was 250 °C, MS Ionization source temperature was 230 °C, the injection port temperature was 250 °C. The samples were injected with 250 split ratio. The injection volume was 1 µL. Oven temperature was programmed in the range of 50 to 250 °C at 3°C/min. The structure of each compound was identified by comparison with their mass spectrum (Nist 05 and Wiley 7 library) [16].

Attenuated Total Reflectance Fourier transform infrared spectroscopy (ATR-FTIR) measurements were run with a Jasco instrument (model 4200), in the following conditions: wavenumber range – 600-4000 cm⁻¹; data points – 3610; aperture setting – 7.1 mm; scanning speed – 2 mm/s; number of scans – 30; resolution – 4 cm⁻¹; filter – 30 kHz; angle of incident radiation – 45° [17].

Total amount of phenolic compounds in the essential oil, at a concentration of 400 ppm in ethanol, was measured with a Jasco V-550 UV-Vis-NIR spectrophotometer. The absorption was measured after 30 min at 20°C, at 765 nm.

Applying essential oil of cloves on leather samples was made by dropping 0.2, 0.4 and 0.6 mL oil and ethanol 1:1 on the surface of 2.0 x 2.0 cm².

The samples treated with essential oil and untreated were placed in each Petri dish in the center of the surface of the culture medium, then the culture medium was seeded in 3 points around the sample, as an equilateral triangle. Petri dishes were placed in thermo-hygrostat at 30°C temperature and were analyzed after 7, 14, 21 and 28 days.

The goal was to monitor the influence of the treatment applied to the sample on mold growth through the mold resistance under simulated contamination, according to no.12697/A 9:2008 „Finished leathers. Mold resistance testing” [18, 19].

Optical microscopy images were captured using a Leica stereomicroscope S8AP0 model with optic fiber cold light source, L2, with three levels of intensity, and magnification 40X [20].

Chemical characteristics of the uncoated leathers were determined according to the following standards: moisture (%) – SR EN ISO 4684:2006; the content extractables (%) – SR EN ISO 4648:2008; the content of chromium oxide (%) – SR EN ISO 5398:2008.

**Results and discussions**

*Identification of compounds in the composition of clove essential oil*

Clove essential oil was analysed using GC-MS [16].

Chromatogram for cloves oil is shown in Figure 1, and identification of compounds in their composition is presented in table 1.

![Chromatogram of organic compounds in the cloves essential oil](Fig.1)

### Table 1

| No. | RT  | Amount, % | Compounds                  |
|-----|-----|-----------|----------------------------|
| 1   | 33.88| 78.03     | Eugenol                    |
| 2   | 36.47| 9.46      | Caryophylene               |
| 3   | 38.01| 1.08      | Alpha Humulene             |
| 4   | 40.58| 10.93     | Aceteugenol                |
| 5   | 43.47| 0.51      | Caryophylene oxide         |
The following compounds are found in the highest amount: eugenol – 78.03%, aceteugenol – 10.93%, caryophylene – 9.46% and alpha humulene – 1.08%.

Clove essential oil was analysed using FT-IR. [17]

FT-IR (ATR) spectra of clove essential oil is shown in Figure 2.

The main bands of clove oil are (Fig. 2): 3509 and 2934 cm⁻¹ – indicating the presence of aliphatic CH₂ groups, 1607 cm⁻¹ and 1761 cm⁻¹ – indicating the presence of C=O group from ester, 1430 and 1368 cm⁻¹ – assigned to the C-H group, 1264 cm⁻¹, 1229 cm⁻¹, 1093 cm⁻¹, 909 cm⁻¹ given by the C-O-C group from ether.

Clove essential oil was analysed using UV-VIS for determination of total amount of phenolic compounds. UV-VIS spectra of clove essential oil is shown in Figure 3.

UV absorption spectra of oil samples in ethanol show the presence of tannins, because the absorption peak observed at 200-280 nm is typical for condensed benzene ring system, in which conjugation is realized between aromatic group and carbonyl group, double bond, or hetero atom.

Characterization by chemical analyses

Chemical characteristics of the uncoated leathers used to obtain bovine hides into natural grain box determined in accordance with standards no. 1619:1994 (Table 2).

Chemical characteristics of the uncoated leathers were determined according to the following standards: moisture (%) – SR EN ISO 4684:2006; the content extractables (%) – SR EN ISO 4648:2008; the content of chromium oxide (%) – SR EN ISO 5398:2008.
Chemical characteristics of the natural grain Box bovine are within the limits specified in standard.

**Biological characterisation of the leather samples**

The samples treated with different amounts of cloves essential oil on the surface of unfinnished leather, CLO-1 (0.2 mL oil and ethanol 1:1), CLO-2 (0.4 mL oil and ethanol 1:1) and CLO-3 (0.6 mL oil and ethanol 1:1), were inoculated with biological material – *Aspergillus niger* spores.

*Aspergillus niger* spores were inoculated in three areas: right side, center and left side of the sample, according to the procedure specified in ASTM D 4576-86 „Standard test method for mold growth resistance of blue stock (leather)”. Incubation was 28 days, but observations were also performed at 7, 14, 21 and 28 days [18, 19].

*Aspergillus niger* strain development was assessed by ranking: 0 – absence of stems and a strong fungitoxic effect, 5 – an almost non-existent effect (the mold covers the entire surface of the specimen). Mold development on leather specimens, and macroscopic images of samples treated with CLO-1-3, after 7, 14, 21 and 28 days from treatment, are presented in table 3.

The numbers under the images are the marks given according to the standard.

**Table 2**

| Sample/Characteristics | CLO-1 | CLO-2 | CLO-3 | M | ST 1619:1994 |
|------------------------|-------|-------|-------|---|--------------|
| Moisture, %            | 14.85 | 14.55 | 14.24 | 14.28 | 14-15        |
| The content extractables, % | 7.12 | 7.68 | 7.39 | 7.87 | Max.8        |
| The content of chromium oxide, % | 5.44 | 5.14 | 5.65 | 5.85 | Min.3.5 |

**Table 3**

| Sample/Characters | 7 | 14 | 21 | 28 |
|------------------|---|----|----|----|
| CLO-1            | ![Image](image1.png) | ![Image](image2.png) | ![Image](image3.png) | ![Image](image4.png) |
| CLO-2            | ![Image](image5.png) | ![Image](image6.png) | ![Image](image7.png) | ![Image](image8.png) |
| CLO-3            | ![Image](image9.png) | ![Image](image10.png) | ![Image](image11.png) | ![Image](image12.png) |
| Control          | ![Image](image13.png) | ![Image](image14.png) | ![Image](image15.png) | ![Image](image16.png) |
The cloves essential oil used to treat box bovine leathers, improves their quality and resistance to fungi and bacteria. Leather samples CLO-1-3 do not develop fungi for 28 days – mark 0). Leather Control sample untreated with the cloves essential oil develops fungi, as shown by marks ranging between 2 after 7 days, 4 after 14 and 5 after 21-28 days.

Conclusions
- Essential oil isolated from cloves (Eugenia caryophyllata) containing: eugenol – 78.03%, aceteugenol – 10.93%, caryophylene – 9.46%.
- The cloves essential oil used to treat box bovine leathers, improves their quality and resistance to fungi and bacteria, reducing the surface defects of natural skin caused by fungi and bacteria.
- The selected essential oil of cloves (due to the high eugenol content) can be used as an antifungal agent in the field of natural leather processing.

Acknowledgements: This works were carried out with the support of Nucleus Program, TEX-PEL-VISION 2022, 4N/8.02.2019/Act ad.l/2019, with the support of the Ministry of Research and Innovation, project number PN 19 17 02 01/2019, and within Program 1 – Development of the national RD system, Subprogram 1.2-Institutional Performance-RDI excellence funding projects, Contract no. 6PFE / 16.10.2018.XS.

References
1. *** Directive 2010/75/EU of the European Parliament and of the Council of 24 November 2010 on industrial emissions (integrated pollution prevention and control).
2. DESELNICU, V., DESELNICU, D.C., VASILESCU, A.M. and MILITARU, G. (2014), “EU Policy for Sustainable Consumption and Production – EU Ecolabel for footwear”, Proceedings of the 5th International Conference on Advanced Materials and Systems ICAMS 2014, 23-25 October 2014, Bucharest, 641-646.
3. DESELNICU, D.C. (2014), European Policies for Products and their Relevance for the Footwear Sector (RO), ISBN: 978-973-720-555-1 (245pg), Agir Press.
4. http://www.europeana.ro/dosare/mediu.htm.legislatie.
5. NZEAKO, B.C., AL-KHAROUSI Z.S.N. and AL-MAHROOQUI, Z., Antimicrobial Activities of Clove and Thyme Extracts, Sultan Qaboos University Medical Journal, 2006, 6(1), 33–39.
6. RADWAN, I.A., ABED, A.H., ABEER, M.R., IBRAHIM, R.A. and ABDALLAH, A.S., “Effect of thyme, clove and cinnamon essential oils on Candida albicans and moulds isolated from different sources”, Mycobiology, 2007, 25(4), 241–243.
7. CHEEN, H.Y. and LEE, M.H., Antifungal Activity of Clove Essential Oil and its Volatile Vapour Against Dermatophytic Fungi, Mycobiology, 2007, 35(4), 241–243.
8. NICULESCU, O., LECI, M., MOLDOVAN, Z., DESELNICU, D.C., Obtaining and characterizing a product with antifungal properties based on essential oils and natural waxes for finishing natural leathers, Rev. Chim. (Bucharest), 66, no. 11, 2015, p. 1733
9. NICULESCU, O., DESELNICU, D.C., GEORGESCU, M., NITURCA, M., Finishing product for improving antifungal properties of leather, Rev. Pielară și Încălzminte (Leather and Footwear J.), 2017, 1, p. 31-38.
10. SIRVAITYTE, J., SIUGZDAITE, J., VALEIKA, V., Application of commercial essential oils of Eucalyptus and Lavender as natural preservative for leather tanning industry, Rev. Chim. (Bucharest), 62, no. 9, 2011, p. 884.
11. European Pharmacopeia, vol. II, ESCOP Strasbourg, Council of Europe, 2005.
12. Farmacopea Română, ediția a X-a, Ed. Medicală, București, 1998.
13. CONSTANTINESCU, D.G., HATIEGANU, E., BUSURICI, F., Plante medicinale utilizate în terapeută, București, Editura Medicală, 2004.
14. STANESCU, U., MIRON, A., HANCIANU, M., APROTOSOAIE, C., Plante medicinale de la A la Z, Iași , Editura Gr. T. Popa, 2004.
15. ARDELEAN, A., MOHAN, G., Flora medicinală a României, București, Editura All, 2008.
16. DAVID, V., MEDVEDOVICI, A., Metode de separare și analiză chromatografică, ediția a II-a, Ed. Universității din București, 2008, 148.
17. MOLDOVAN, Z., Metode instrumentale de analiză, Ed. Universității din București, 2001.
18. ST 12697/A 91/2008, standard for determination of leather resistance to fungi.
19. ST 21576/A 91/2008, standard test method for mold growth resistance of blue stock (leather).
20. XXX-Manual de utilizare a microscopului optic.

Manuscript received: 13.03.2019