

Abstract



Extraction and HPLC-UV Analysis of Usnic Acid in Usnea barbata Collected in the Altai Krai⁺

Georgiy Kutateladze 1,* and Irina Anikina 2

3 4

5

6

7

8 9

10

28

29

30

39

1

2

- ¹ Department of Pharmacy, Altai State Medical University, Ministry of Health of Russian Federation, 40, Lenina Str., 656038 Barnaul, Russia; gohakutateladze@gmail.com
- ² Department of Pharmacy, Altai State Medical University, Ministry of Health of Russian Federation, 40, Lenina Str., 656038 Barnaul, Russia; anikina_bri@mail.ru
- * Correspondence: gohakutateladze@gmail.com
- + Presented at the title, place, and date.

Abstract: Usnea barbata L. (Parmeliaceae Zenker.) is used in folk medicine as an antimicrobial agent 11 in various diseases of bacterial genesis. The antimicrobial activity is associated with the presence of 12 the lichen acids, especially usnic acid (UA). Aim of the study is isolation, identification and assay of 13 UA by HPLC-UV. The lichen Usnea barbata was collected from tree branches in the Altai Krai during 14 the growing season (June-July 2022). The sum of lichen acids was isolated in the form of sodium 15 salts by re-extraction with 20% sodium hydroxide solution from chloroform extract. The re-extract 16 was acidified with 10% hydrochloric acid until a precipitate formed, which was filtered off, dried, 17 dissolved in acetonitrile and analyzed. The HPLC analysis was conducted using a MiliChrom A-02 18 HPLC System equipped with UV-spectrophotometric detector. The chromatographic conditions: 19 column - reverse-phase, ProntoSIL 120-5 C18 (75×2.0 mm, 5 µm); column oven temperature - 40°C; 20 mobile phase (MP) - 0.1% trifluoroacetic acid aqueous solution (solvent A), 100% acetonitrile (sol-21 vent B); MP flow rate - 150 µL/min with gradient elution - from 10 to 50% of solvent B in 5 minutes, 22 gradually increased to 100% up to 20 minutes; sample injection volume - 4 µL; detection at 230 and 23 280 nm. By comparing the retention times ($\tau = 15.0\pm0.1$ min) and spectral characteristics (λ max) with 24 those of the 0.05% UA reference sample solution, the presence of UA in the studied thallus was 25 established. Quantitative content, calculated from the peak area, compared with the peak area of 26 the reference sample -1.45±0.07%. 27

Keywords: *Usnea barbata*; usnic acid; HPLC-UV

Supplementary Materials:

Author Contributions: Conceptualization, G.K. and I.A.; methodology, G.K.; formal analysis, G.K.;31investigation, G.K.; resources, G.K.; data curation, G.K.; writing—original draft preparation, G.K.;32writing—review and editing, I.A.; visualization, G.K.; project administration, G.K. All authors have33read and agreed to the published version of the manuscript.34

- Funding: This research received no external funding.35Institutional Review Board Statement: Not applicable.36Informed Consent Statement: Not applicable.37Data Availability Statement: Not applicable.38
 - **Conflicts of Interest:** The authors declare no conflict of interest.

Citation: Kutateladze, G. Title. Med. Sci. Forum 2023, 2, x. https://doi.org/10.3390/xxxxx

Academic Editor: Firstname Lastname

Published: date

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2023 by the authors. Submitted for possible open access publication under the terms and conditions of the Creative Commons Attribution (CC BY) license (https://creativecommons.org/licens es/by/4.0/).