

Spectroscopy. NMR spectra were determined at 60 MHz, in CCl_4 and with trimethylsilane as internal standard; values are given in ppm (δ) relative to trimethylsilane.

Catalytic hydrogenation of the norseychellane. Catalytic hydrogenation of the norcapillene (55 mg) in EtOH (5 ml) over PtO_2 (1.5 mg) was carried out at room temp. for 2.5 hr. The product was purified by preparative TLC (kieselgel GF254) using *n*-hexane C_6H_6 (2:1), as was obtained as a colorless oil. NMR: δ^{CCl_4} 0.88 (3H, t, $\text{CH}_2\text{-CH}_3$), 1.26 (6H, m, $-(\text{CH}_2)_3-$), 2.68 (2H, t, $\phi\text{-CH}_2-$), 7.23 (5H, m, ϕ). (Found: C, 89.11%; H, 10.89%. Calcd for $\text{C}_{11}\text{H}_{16}$: C, 89.12%; H, 10.88%).

REFERENCES

1. Miyazawa, M. and Kameoka, H. (1975) *Phytochemistry* **14**, 1126.
2. Miyazawa, M. and Kameoka, H. (1975) *Phytochemistry* **14**, 1126.
3. Quang, L. V. and Cadiot, P. (1965) *Bull. Soc. Chem. France* 1525.
4. Taniguchi, H., Mathai, I. M. and Miller, S. I. (1966) *Tetrahedron* **22**, 867.
5. Quang, Y. V., Quang, L. V. and Emptoz, G. (1964) *Compt. Rend.* **258**, 4586.
6. Migniac, L. G. (1961) *Ann. Chim.* **6**, 1071.

Phytochemistry, 1976, Vol. 15, p. 224. Pergamon Press. Printed in England.

NORSEYCHELANONE, α - AND β -PATCHOULENES AND PATCHOULI ALCOHOL FROM *NARDOSTACHYS JATAMANSI*

G. RÜCKER*, J. TAUTGES*, M. L. MAHESWARIT† and D. B. SAXENA†

* Institut für Pharmazeutische Chemie der Westfälischen Wilhelms-Universität Münster, BRD;

† Division of Agricultural Chemicals, Indian Agricultural Research Institute, New Delhi 110012, India

(Received 24 June 1975)

Key Word Index—*Nardostachys jatamansi*; Valerianaceae; norseychelanonone; α - and β -patchoulenes; patchouli alcohol.

Plant Nardostachys jatamansi DC, Valerianaceae. **Origin:** India, from the Himalaya mountains at a height of 3000–5000 m. The plant purchased from the local market. **Previous work** on the roots [1–3].

Isolation 24 kg air-dried roots are exhaustively extracted with light petrol (35–40°). Solvent was evaporated *in vacuo* and the residue (900g) extracted with Na_2CO_3 -solution. The insoluble portion was chromatographed on a silica gel column (180 $\times$ 2.0 cm) using light petroleum (60°)–EtOAc, 8:2 as eluent. The fraction with R_f 0.78 (TLC same solvent) was found to be norseychelanonone isolated for the first time from any plant source. The fraction R_f 0.70, formed colourless crystals mp 55–56 (patchouli alcohol). The fractions having R_f 0.90–1.0 were rechromatographed on silica gel columns impregnated with 10% AgNO_3 , using hexane as an eluent. TLC showed spots at R_f 0.5, 0.60, 0.68 and 0.80. The bright red spot (R_f 0.45) was identified as seychellen [4,5], confirmed by IR, NMR and MS.

Identification Norseychelanonone: Identical with the compound obtained by degradation of seychellen, TLC, IR, NMR, MS [4,5]. **Patchouli alcohol:** IR [6–9], MS [7–9], TLC (Kieselgel Merck PF₂₅₄), Anisaldehyde– H_2SO_4 : carmine red, R_f : 0.70. α - and β -Patchoulenes: by GC-MS (Varian-CH7), 3.8% SE 30, 100°C/6' 2 min to 250°. The GLC of the mixture showed 4 peaks having R_t : 5.3/9.15/10.0 and 11.0 min. For further identification by GLC 50% reference substances were added to the mixture. β -Patchoulene showed an increase in the height of the peak

with R_t 9.15 min. MS: $m/e(\%)$ 204(60M⁺), 189(98), 161(100), 156(30), 135(30), 133(40), 119(70), 105(60), 93(50), 91(50), 79(30), 77(25). α -Patchoulene showed an increase in the height of the peak R_t 11.0 min. MS: $m/e(\%)$ 204(20M⁺), 189(20), 161(30), 135(80), 119(45), 108(50), 107(90), 105(55), 93(100), 91(50), 81(20), 79(30), 77(30), 55(50).

Acknowledgements—We thank Dr. Bruhn and Dr. Klein, DRAGOCO Co., Holzminden, BRD, for supplying the spectra and reference substances.

REFERENCES

1. Rücker, G. and Glauch, G. (1967) *Deut. Apotheker-Z.* **27**, 921.
2. Sastry, S. D., Maheswari, M. L., Chakravarti, K. K. and Bhattacharyya, S. C. (1967) *Ess. Oil Record* **58**, 154.
3. Rücker, G. and Tautges, J. (1974) *Arch. Pharmaz.* **307**, 791.
4. Wolf, G. and Ourisson, G. (1969) *Tetrahedron* **25**, 4903.
5. Maheswari, M. L. and Saxena, D. B. (1974) *Ind. J. Chem.* **12**, 1221.
6. Büchi, G. and Erickson, R. N. (1956) *J. Am. Chem. Soc.* **78**, 1262.
7. Yamaguchi, K. (1970) *Spectral Data of Natural Products* Vol. 1. Elsevier, Amsterdam.
8. Hill, H. C., Reed, J. R. and Robert-Lopes, M. T. (1968) *J. Chem. Soc. C (London)* p. 93.
9. Dobler, M., Dunitz, J. D., Gubler, B., Weber, H. P., Büchi, G. and Padilla, J. (1963) *Proc. Chem. Soc. (London)* p. 383.