

min the mixture was centrifuged to remove inorganic matter, solvent was removed *in vacuo* and the residue crystallized from EtOAc-petrol (1:2) to give the quinone (5) mp 62–64° (sublimes at 58°).

Acknowledgements—This work was supported by a grant, AI-00226 from the National Institute of Allergy and Infectious Diseases, and a grant RRO5621, from the Division of Research Resources, National Institutes of Health. We thank Mr. I. Miura for determining several of the NMR spectra. The authors also thank Mr. Francis Manginelli and Mr. Thomas Amato for capable technical assistance.

REFERENCES

1. Nair, M. S. R. and Anchel, M. (1975) *Tetrahedron Letters* **31**, 2641.
2. Jackman, L. M. and Sternhell, S. (1969) *Applications of Nuclear Magnetic Resonance Spectroscopy in Organic Chemistry*, p. 136. Pergamon Press, New York.
3. Ishibashi, K., Nose, K., Shindo, T., Arai, M. and Mishima, H. (1968) *Ann. Sankyo Res. Labs.* **20**, 76.
4. Levy, G. C. and Nelson, G. L. (1972) *Carbon 13 Nuclear Magnetic Resonance for Organic Chemists*, p. 71. Wiley-Interscience, New York.
5. Farrel, I. W., Keeping, J. W., Pellatt, M. G. and Thaller, V. (1973) *J. Chem. Soc. Perkin Trans.* 2642.
6. Turner, W. B. (1971) *Fungal Metabolites*, p. 41. Academic Press, New York.
7. Ballogh, V., Fetizon, M. and Golfier, M. (1971) *J. Org. Chem.* **36**, 1339.

ISOLATION OF SOLAVETIVONE FROM *NICOTIANA TABACUM*

TAKANE FUJIMORI, REIKO KASUGA, HAJIME KANEKO and MASAO NOGUCHI

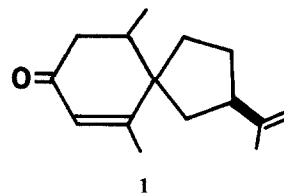
Central Research Institute, The Japan Tobacco and Salt Public Corporation, Umeoka 6-2, Midori-ku, Yokohama, 227, Japan

(Received 31 August 1976)

Key Word Index—*Nicotiana tabacum*; Solanaceae; Burley tobacco; vetispirane derivative; solavetivone

Preparative GLC of the neutral oil from 370 kg of air-cured American Burley tobacco leaves [1] afforded 10 mg colourless mobile oil (1) $[\alpha]_D^{23} - 92.8^\circ$ (EtOH; c 0.15). The MS spectrum showed significant peaks at m/e 218(M^+ , 48%), 137(58), 133(64), 108(100), 93(86), 79(72), 68(70), 67(70) and 41(74). By the high resolution MS, the formula of (1) was estimated as $C_{15}H_{22}O$ (found: 218.1676, calcd.: 218.1670). The IR spectrum showed an α, β -unsaturated carbonyl group at 1669 cm^{-1} and a terminal methylene group at 3090, 1650 and 893 cm^{-1} . No hydroxyl group was observed. The NMR spectrum (100 MHz, $CDCl_3$) showed a secondary methyl group (δ 0.98, d , 7.0 Hz, 3H), a methyl group attached to a β -position of α, β -unsaturated carbonyl group (δ 1.93, d , 11.2 Hz, 3H), an isopropenyl group (δ 1.75, $br\ s$, 3H and δ 4.68, $br\ s$, 2H), a methylene group adjacent to a carbonyl group (δ 2.55, AB part of an ABX system) and an olefinic methine (δ 5.62, $br\ s$, 1H). The UV spectrum had λ_{max}^{EtOH} 241.5 nm (ϵ 14000) which was consistent with an enone system. These spectroscopic data of (1) were identical with those of solavetivone, 6,10-dimethyl-2-(1-methylethenyl)-spiro[4.5]dec-6-en-8-one, which has been isolated as a major stress metabolite produced by infected potato tubers [2]. This compound was also

present in the leaves of *N. tabacum* cv Matsukawa, one of representative Japanese domestic tobacco, and Phillipine cigar tobacco (Manila leaves). This is the first report of the naturally existence of a vetispirane derivative in *N. tabacum*.



Acknowledgement—The authors are indebted to Mr. Saizo Shibata of this laboratory for high resolution mass spectrometric analysis.

REFERENCES

1. Fujimori, T., Kasuga, R., Matsushita, H., Kaneko, H. and Noguchi, M. (1976) *Agr. Biol. Chem.* **40**, 303.
2. Coxon, D. T., Price, K. R., Howard, B., Osman, S. F., Kalan, E. B. and Zacharius, R. M. (1974) *Tetrahedron Letters* **34**, 2921.