

M.p., mixed m.p., IR, NMR. *Glycerides of fatty acids*. Acid part consists of C₂₄–C₃₂ straight-chain saturated fatty acids. Identified by GLC of methyl esters. *Unidentified compound*. (A) C₃₀H₅₀O, m.p. ~50°, IR ν^{KBr} 3400, 1030, 812 cm⁻¹, NMR $\delta_{\text{CDCl}_3}^{\text{TMS}}$ 0.75 (3H,s), 0.81 (3H,s), 0.86 (3H,s), 0.88 (3H,d, $J = 6.3$), 0.97 (6H,s), 1.60 (3H,s), 1.68 (3H,s), 3.25 (2H,m), 5.09 (1H,m), 5.25 (1H,m), benzoate, m.p. 144–145°, IR.

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LABIATAE

ESSENTIAL OIL FROM THE LEAVES AND INFLORESCENCE OF *OCIMUM GRATISSIMUM*

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Plant. Ocimum gratissimum L. Indigenous to Nigeria.

Previous studies. Major component of oil from specimens collected in Taiwan has been shown¹ to be eugenol (62%). This compound is the major component of the oil obtained from the leaves of a hybrid between *O. gratissimum* and *O. menthaefolium*.² The Nigerian *O. gratissimum* was reported^{3,4} to contain thymol, but no eugenol. The remaining components of the oil from the Nigerian plant are reported here.⁵

RESULTS

Composition of oil from leaves (%):— α -pinene (2.6), camphene (4.0), β -pinene (0.6), α -terpinene: Δ^3 -carene (4.1), myrcene (1.4), 1,8-cineole (1.1), α -terpinene (6.2), p -cymene (16.2) limonene (1.8) camphor (0.6), linalool (0.2), α -terpineol (2.4), C₁₀H₂₂O (2.3), thymol (47.6), methyleugenol (1.7), methylisoeugenol (trace), caryophyllene (2.1), humulene (0.5), β -selinene (1.6), longifoline (3.0), clovene (trace). Oil from the flowers has essentially the same composition except the proportion of camphene is reduced.

¹ PING-HSIEN YEH, *Perfumery Essent. Oil Record*, **51**, 611 (1960).

² O. K. MADALSKA, C. BANKOWSKI and J. KUDUK, *Acta. Polon. Pharm.* **21**, 387 (1964).

³ F. EL-SAID, E. A. SOFOWORA, S. A. MALCOLM and A. HOFER, *Planta Med.* **17**, 195 (1969).

⁴ E. A. SOFOWORA, *Planta Med.* **18**, 173 (1970).

⁵ M. QUDRAT-I-KHUDA, M. ERFAN ALI, A. KHALIQUE and L. A. M. SHAMSUZZAMAN, *Sci. Res. (Dacca, Pakistan)* **1**, 217 (1964).

EXPERIMENTAL

The oils were obtained by steam distillation of freshly gathered leaves and flowers collected at the onset of flowering. Analysis was by GLC and GLC-MS (Column conditions: 15% carbowax 20 M on chromosorb W, 15% Apiezon L. on Universal B and 5% SE 30 gum rubber on Universal B. *Instrumental*. Pye 104 gas chromatograph coupled to MS 12 Mass-spectrometer). Individual compounds were characterized by direct comparison with the retention indices and fragmentation patterns of authentic specimens.

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LAURACEAE

n-PARAFFINS FROM THE LEAVES OF THREE GENERA OF LAURACEAE

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Abstract—*n*-Paraffins ranging from C₁₆ to C₃₃ were detected by gas chromatography from the leaves of *Cinnamomum camphora* Sieb., *Lindera obtusiloba* Blume, *Litsea japonica* Juss.).

Plants. *Cinnamomum camphora* Sieb., *Lindera obtusiloba* Blume, *Litsea japonica* Juss.

Occurrence. Hiroshima Prefecture, Japan.

Previous work. No work (concerning paraffin constituents).

Date. *Cinnamomum camphora* Sieb. (January 1971), *Lindera obtusiloba* Blume and *Litsea japonica* Juss. (September 1970).

Leaves. Crushed to pieces and extracted with *n*-hexane. Purification (column chromatography and molecular sieve 5A treatment).¹ Identification (GLC using two columns, SE-30-5%, Apiezon grease L-5%, column temperature 150–300°). The odd paraffins are in large amount (*Cinnamomum*, *Lindera*, *Litsea* are 84.2, 88.6, 57.9%, even ones are 13.8, 11.4, 42.1% respectively).

n-Paraffins. *Cinnamomum camphora* Sieb.: C_{16–19} (trace), C₂₀(0.2%), C₂₁(0.5), C₂₂(0.8), C₂₃(2.6), C₂₄(2.4), C₂₅(7.1), C₂₇(23.9), C₂₈(4.3), C₂₉(43.3), C₃₀(1.2), C₃₁(8.8), C_{32–33}(trace), *Lindera obtusiloba* Blume: C₁₆(1.1), C₁₇(0.3), C₁₈(1.3), C₁₉(0.4), C₂₀(1.3), C₂₁(0.4), C₂₂(1.3), C₂₃(1.4), C₂₄(1.3), C₂₅(3.9), C₂₆(1.9), C₂₇(25.6), C₂₈(1.6), C₂₉(45.1), C₃₀(1.6), C₃₁(11.5), C_{32–33}(trace). *Litsea japonica* Juss.: C_{19–22}(trace). C₂₃(0.6), C₂₄(2.1), C₂₅(3.4), C₂₆(7.0), C₂₇(14.4), C₂₈(18.8), C₂₉(23.8), C₃₀(14.2), C₃₁(14.8), C₃₂(trace), C₃₃(0.9).

¹ N. Y. CHEN and S. J. LUCKI, *Analyt. Chem.* **42**, 508 (1970).