



RJC

http://www.rasayanjournal.com

RASĀYAN J. Chem.
Vol.2, No.1 (2009), 156-158
ISSN: 0974-1496
CODEN: RJCABP

SHORT COMMUNICATION

CHEMICAL CONSTITUENTS OF *ARTABOTRYS*
ODORATISSIMUS (SEEDS)

Jagdish P. Singh^{*1}, Alok K. Singh¹, Archana Singh¹ and Rakesh Ranjan²

¹Department of Chemistry, S.G.R.-Post Graduation College, Dobhi, Jaunpur-222149, India.

²Department of Medicinal Chemistry, Institute of Medical Sciences,
Banaras Hindu University, Varanasi-221005, India.

*E-mail: sinharanjan1@gmail.com

ABSTRACT

3-Hydroxy-9-methoxypterocarpan, nonacosanoic acid 2', 3'-dihydroxypropyl ester, pentacosanoic acid 2,-3'-dihydroxypropyl ester and docosanoic acid have been isolated for the first time from the seeds of *Artabotrys odoratissimus* and identified by spectroscopic data of these natural products and their derivatives.

Key words: 3-Hydroxy-9-methoxypterocarpan and cosanoic acids derivative.

INTRODUCTION

Artabotrys odoratissimus R. Brown, belongs to the family annonaceae which is an evergreen perennial shrub, mostly native of tropical Africa, Eastern Asia and India¹. This plant is used in Indian system of medicine for the treatment of vomiting, biliousness and disease of blood and heart. The leaves are reported to contain an anti-fertility principal². Oil of seeds of the plants are reported to contain antidandruff, anti-itching, anti-thrombin and hair loss preventing activities³. In continuation to our previous communication⁴, we report here in isolation and identification of four compounds viz; 3-Hydroxy-9-methoxypterocarpan, nonacosanoic acid 2', 3'-dihydroxypropyl ester, pentacosanoic acid 2', 3'-dihydroxypropyl ester and docosanoic for the first time from this plant.

EXPERIMENTAL

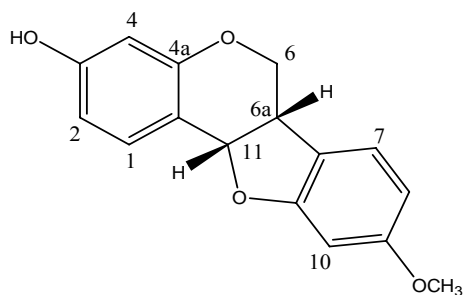
The melting points were measured on a Yazawa hot stage microstage apparatus and are uncorrected. Optical rotations were measured on JASCO DIP-360 Polarimeter (Cell length 5 CM). UV absorption spectra were recorded on JASCO UV/visible spectrophotometer (model no. 7800) while IR on JASCO FT-IR 5300 spectrometer. NMR spectra were recorded on an AMX 300 NMR spectrometer (Bruker, Karlsruhe, Germany).

The seeds of *A. odoratissimus* were collected from the garden of Banaras Hindu University and then authenticated by Professor. V. K. Joshi, Department of Dravya Guna, IMS, Banaras Hindu University, Varanasi-221005, INDIA.

RESULTS AND DISCUSSION

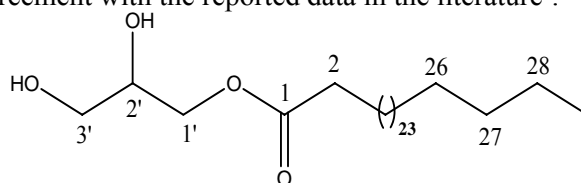
The milled seeds (1 kg) were extracted with 95% ethanol by cold percolation method, solvent was removed under vacuum at 40°C. The left residue (80 gm) was then fractionated into *n*-hexane, chloroform, *n*-butanol and aqueous soluble fraction. The chloroform soluble fraction was chromatographed over silica gel. Elution of column with solvents of increasing polarity resulted in the isolation of four compound viz; 3-hydroxy-9-methoxypterocarpan (25 mg, *n*-hexane: EtOAc 1:1), nonacosanoic acid 2', 3'-dihydroxypropyl ester (22mg, EtOAc), pentacosanoic acid 2,-3'-dihydroxypropyl ester (24mg, EtOAc : MeOH 9:1, early fraction) and docosanoic acid(20 mg, EtOAc : MeOH 9:1, latter fraction).

Compound 1. $[\alpha]_D^{22} -226^0$ (c, 0.1, CHCl_3); MF: $\text{C}_{16}\text{H}_{14}\text{O}_4$, M^+ at m/z 270 (EI-MS); *m.p.* 127-128 ^0C ; IR (KBr) ν_{max} 3404, 2949, 1619, 1600, 1499, 1474, 1365, 1281, 1149, 1028, 934, 837, 764 cm^{-1} ; UV λ_{max} 282, 287 (MeOH) nm; $^1\text{H-NMR}$ (CDCl_3): δ_{H} 7.37 (1H, *d*, $J=8.3$, H-1), 6.54 (1H, *d*, $J=8.3$, H-2), 6.44-6.41 (*m*, H-4), 3.67-3.51 (*m*), 4.23 (1H, *dd*, $J=6.0, 9.8$, H-6), 3.67-3.51 (*m*, H-6a), 7.12 (1H, *d*, $J=8.7$, H-7), 6.56-6.41 (H-8 & H-10), 5.49 (1H, *d*, $J=6.0$), 3.76 (3H, *s*, 9-OCH₃), 5.59 (*br hump*, 3-OH); $^{13}\text{C-NMR}$: δ_{C} 132.4 (C-1), 110.1 (C-2), 157.5 (C-3), 103.8 (C-4), 156.8 (C-4a), 66.7 (C-6), 39.7 (C-6a), 119.4 (C-6b), 125.0 (C-7), 106.6 (C-8), 161.2 (C-9), 97.1 (C-10), 160.8 (C-10a), 78.8 (C-11a), 112.6 (C-11b), 55.7 (-OCH₃). On the basis of above spectral data the structure of this compound was settled as 3-hydroxy-9-methoxypetrocarpans commonly known as (-) medicarpen. The above spectral values were in good agreement with the reported data in the literature^{5,6,7}.



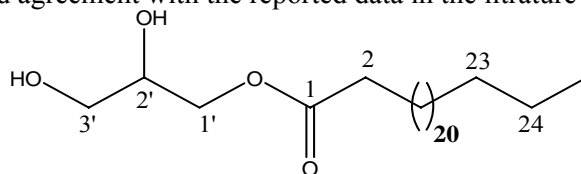
Compound-1

Compound 2. $[\alpha]_D^{22} -3.87$ (MeOH + CHCl_3 , c, 0.10); MF: $\text{C}_{32}\text{H}_{64}\text{O}_4$, M^+ at m/z 512 (EI-MS); *m.p.* 90-91 ^0C ; IR (KBr) ν_{max} 3425, 2919, 2851, 2363, 1734, 1468, 1179, 1051, 720 cm^{-1} ; $^1\text{H-NMR}$ (CDCl_3 +DMSO- d_6 , 200 MHz): δ_{H} 4.04 (2H, *t*, $J=5.3$ Hz, H-1'), 3.80 (1H, *m*, H-2'), 3.48 (2H, *m*, H-3'), 2.29 (2H, *t*, $J=7.3$ Hz, H-2), 1.56 (6H, *brm*, H-26,27,28), 1.24 (46H, *brs*, H-3 to H25), 0.87 (3H, *t*, $J=6.0$ Hz, CH₃); $^{13}\text{C-NMR}$ δ_{C} 172.6 (C-1), 68.9 (2'), 64.3 (C-1'), 62.3 (C-3'), 33.1 (C-2), 30.7 (C-28), 28.5 (C-7 to C-27), 28.2 (C-3), 28.0 (C-4), 23.8 (C-5), 21.5 (C-6), 13.1 (CH₃). On the basis of above spectral data the structure of compound 2 was established as nonacosanoic acid 2', 3'-hydroxy-propyl ester. The above spectral data are in good agreement with the reported data in the literature⁸.



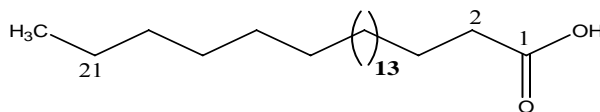
Compound-2

Compound 3. $[\alpha]_D^{22} -3.11$ (MeOH+ CHCl_3 , c, 0.22); MF: $\text{C}_{28}\text{H}_{56}\text{O}_4$, FAB-MS: m/z 457 $[\text{M}+\text{H}]^+$; *m.p.* 72-74 ^0C ; IR (KBr) ν_{max} 3225, 1733, 1704, 1389, 725 cm^{-1} ; $^1\text{H-NMR}$ (CDCl_3 + DMSO- d_6): δ_{H} 4.11 (2H, *t*, $J=6.2$ Hz, H-1'), 3.80 (1H, *m*, H-2'), 3.34 (2H, *m*, H-3'), 2.26 (2H, *t*, $J=7.3$ Hz, H-2), 1.56 (4H, H-23,24), 1.25 (40H, *brs*, H-3 to H-22), 0.87 (3H, *t*, $J=6.0$ Hz, CH₃); $^{13}\text{C-NMR}$ δ_{C} 172.2 (C-1), 69.0 (2'), 66.5 (C-1'), 64.5 (C-3'), 32.9 (C-2), 30.7 (C-24), 30.3 (C-6 to C-23), 30.1 (C-3), 25.9 (C-4), 23.7 (C-5), 14.1 (CH₃). Thus, the structure of compound 3 was established as pentacosanoic acid 2, 3-dihydroxypropylester which is in good agreement with the reported data in the literature⁹.



Compound-3

Compound 4. MF: C₂₂H₄₄O₂, M⁺ at m/z 340; m.p. 70-72⁰C; IR (KBr) ν_{\max} 3430, 2919, 2850, 2362, 1693, 1594, 1468, 1351 cm⁻¹; 325, 311, 297, 283, 269, 255, 241, 227, 213, 199, 185, 171, 157, 143, 129; ¹H-NMR δ_{H} 2.28, (2H, *t*, J= 7.19 Hz, H₂-2), 1.60 (2H, *m*, H-3), 1.25 (36H, *brs*, H-4 to H-21), 0.87 (3H, *t*, J=6.2 Hz, H-22); ¹³C-NMR δ_{C} 176.1 (C-1), 34.3 (C-2), 25.0 (C-3), 22.7 (C-21), 14.2 (C-22), 29.4 (C-4 to C-20) has suggested it to be docosanoic acid which is consistent with the ¹³C-NMR data of compound 4¹⁰.



Compound-4

ACKNOWLEDGEMENT

R. R. is thankful to UGC for financial assistance.

REFERENCES

1. The Wealth of India: Raw Materials, A dictionary of Indian raw Material and Industrial Products, Council of Scientific Industrial Research, New Delhi, 119 (1952).
2. A.P. Singh and M. Sahai, *Planta Medica.*, **62**, 192 (1996).
3. M. Qata, H. Komazaki, K. Kobayashi, H. C. Sakamoto, C. Hanzawa, Y. Tsuji, T. Mangara and T. Nishi, *Kokai Tokkoyokoho*, **9**, 433 (1997).
4. J.P. Singh, A.K. Singh, A. Singh and R. Ranjan, *Indian Journal of Natural Prods.*, (in press)
5. S.H. Harper, A.D. Kemp and W.G.E. Underwood, *Chem. Ind.*, **562** (1965).
6. G.J.H. Rall and M.E. Oberholzer, *Tetrahedron*, **33**, 1735 (1977).
7. T.B.H. McMurry, E. Martin, D.M.X. Donnelly and J.C. Thompson, *Phytochem.*, **11**, 3283 (1972).
8. E. Poloucek, J. Macicek, J. Petruj, K. Vesely, D. Rysavy, F. Kratochvil, T. Sverak, M. Kummer, V. Douda, and M. Vodicka, Czech Patent CS 247458, 5pp, 1984. *Chem. Abstr.*, **109**, P74562a (1988).
9. Y. Sang, H. Shi and Z. Min, *Zhongcaoyao Zazhi Bianjibu*, **33**, 776 (2002).
10. B. Holbom, *J. Am. Oil Chem. Soc.*, **54**, 289 (1977)

(Received: 20 December 2008

Accepted: 28 December 2008

RJC-313)

"I like to see a man proud of the place in which he lives. I like to see a man live so that his place will be proud of him."

- Abraham Lincoln