#### SHORT COMMUNICATION

## FLAVONOIDS OF THREE CROTALARIA SPECIES

#### S. Sankara Subramanian and S. Nagarajan

Department of Chemistry, Jawaharlal Institute of Postgraduate Medical Education and Research, Pondicherry-6, India

(Received 3 April 1970)

Abstract—The flavonoids of Crotalaria striata, C. paniculata and C. anagyroides have been studied, and apigenin and its glycosides are commonly found.

Plant. Species of Crotalaria (Leguminosae—sub-family Lotoideae).

Uses, Medicinal.1-4

*Previous work.* On sister species. 5-9

#### Present Work

Alcoholic extract of the material fractionated with petrol, ether and EtOAc.

### C. striata DC. (= C. mucronata Desv.)

Leaves. Vitexin (from EtOAc fraction, 0.05% yield, m.p. and mixed m.p., hydrolytic fission with HI in phenol,  $R_f$  and co-chromatography) and vitexin-4'-O-xyloside (from the aq. fraction after EtOAc, by the lead salt method) ( $R_f$  and co-chromatography, hydrolysis to vitexin and xylose).

#### C. striata DC

Stem bark. Apigenin (from ether fraction, acetate, m.p. and mixed m.p.,  $R_f$  and co-chromatography). Vitexin and its 4'-O-xyloside (from EtOAc fraction, confirmed as above).

#### C. paniculata Willd

Flowers. Quercetin 3-galactoside (from EtOAc fraction, m.p. and mixed m.p.,  $R_f$  and co-chromatography, acid hydrolysis to quercetin and galactose), vitexin-4'-Q-xyloside (from EtOAc fraction, confirmed as under C. striata leaves).

- <sup>1</sup> R. N. CHOPRA, S. L. NAYAR and I. C. CHOPRA, Glossary of Indian Medicinal Plants, p. 81, Council of Scientific & Industrial Research, New Delhi (1956).
- <sup>2</sup> K. M. NADKARNI, Indian Materia Medica, Vol. I, p. 391, Popular Book Depot, Bombay (1954).
- <sup>3</sup> K. R. KIRTIKAR and B. D. BASU, *Indian Medicinal Plants* (edited by L. M. BASU), Vol. I, p. 693, Allahabad (1933).
- <sup>4</sup> J. M. WATT and M. G. Breyer-Brandwijk, *The Medicinal and Poisonous Plants of Southern and Eastern Africa*, p. 577, E. & S. Livingstone, London (1962).
- <sup>5</sup> S. Sankara Subramanian and S. Nagarajan, Current Sci., India 36, 364 (1967).
- <sup>6</sup> S. Sankara Subramanian and S. Nagarajan, *Planta Med.* 16, 432 (1968).
- <sup>7</sup> S. Sankara Subramanian and S. Nagarajan, Current Sci., India 36, 403 (1967).
- <sup>8</sup> S. SANKARA SUBRAMANIAN and S. NAGARAJAN, Indian J. Pharm. 29, 311 (1967).
- 9 S. SANKARA SUBRAMANIAN and S. NAGARAJAN, Current Sci., India 38, 65 (1969).

C. anagyroides H. B. and K.

Stem bark. Apiin (from EtOAc fraction, m.p.,  $R_f$  and hydrolysis by 10% H<sub>2</sub>SO<sub>4</sub> to apigenin, glucose and apiose).

C. juncea L.

Seeds. Apigenin-7-glucuronide and apigenin-7,4'-O-di-glucoside (from EtOAc fraction,  $R_f$ ).

Acknowledgement—We thank the Principal, J.I.P.M.E.R., for kind encouragement.

#### SHORT COMMUNICATION

# CHLOROGENIN AND KAEMPFEROL GLYCOSIDES FROM THE FLOWERS OF AGAVE AMERICANA

S. SANKARA SUBRAMANIAN and A. G. R. NAIR

Department of Chemistry, Jawaharlal Institute of Postgraduate Medical Education and Research, Pondicherry-6, India

(Received 3 April 1970)

Abstract—Chlorogenin was isolated in a yield of 0.5% from the fresh flowers of Agave americana. The flavonol glycosides were identified as kaempferol-3-glucoside and kaempferol-3-rutinoside.

Plant. Agave americana L.—Amaryllidaceae.

Source. Pondicherry.

Uses. Medicinal.1,2

Previous work. Hecogenin from leaves; 2,3 work on sister species.4

Present work. Examination of flowers.

Fresh flowers extracted with hot ethanol (95%) under reflux, aq. concentrate shaken and layered with an equal volume of benzene and kept in an ice-chest for 2 weeks. The colourless solid separated at the interphase on crystallization thrice from MeOH yielded chlorogenin,  $^5$   $C_{27}H_{44}O_4$ , m.p. 272-274°,  $[\alpha]_2^{28} - 51\cdot2^\circ$  (py); diacetyl, m.p. 154-155°,  $[\alpha]_2^{28} - 36\cdot5^\circ$ ; dibenzoyl, m.p. 200-203°,  $[\alpha]_2^{28} - 9\cdot8^\circ$ . Benzene concentrate yielded a small quantity of the same solid (total yield,  $0\cdot5^\circ$ ). No hecogenin could be identified. Ether extract of the aq. alc. concentrate yielded small quantity of kaempferol ( $R_f$  and co-chromatography). EtOAc extract yielded two glycosides of kaempferol (separated by preparative PC) identified as kaempferol-3-glucoside and kaempferol-3-rutinoside (m.p.,  $R_f$ , acid hydrolysis and co-chromatography with authentic samples) (total yield of flavonols,  $0.03^\circ$ ).

Acknowledgement-Our thanks are due to the Principal, J.I.P.M.E.R., for encouragement.

<sup>&</sup>lt;sup>1</sup> R. N. CHOPRA, I. C. CHOPRA, K. L. HANDA and L. D. KAPUR, *Chopra's Indigenous Drugs of India*, p. 577, U. N. Dhur, Calcutta (1958).

<sup>&</sup>lt;sup>2</sup> J. M. WATT and M. G. Breyer-Brandwijk, *The Medicinal and Poisonous Plants of Southern and Eastern Africa*, p. 19, E. & S. Livingstone, London (1962).

<sup>&</sup>lt;sup>3</sup> H. SINGH and W. PEREIRA, JR., Indian J. Chem. 2, 297 (1964).

<sup>&</sup>lt;sup>4</sup> K. Paech and M. V. Tracey, *Modern Methods of Plant Analysis*, Vol. III, pp. 191-200, Springer-Verlag, Berlin (1955).

<sup>&</sup>lt;sup>5</sup> G. HARRIS, Dictionary of Organic Compounds, Vol. III, p. 629, Eyre & Spottiswoode, London (1965).