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°,  $[\alpha]_D^{28} - 51\cdot2^{\circ}$  (py); diacetyl, m.p. 154–155°,  $[\alpha]_D^{28} - 36\cdot5^{\circ}$ ; dibenzoyl, m.p. 200–203°,  $[\alpha]_D^{28} - 9\cdot8^{\circ}$ . 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).