

cystidia (figure 2F) extending beyond the hymenial layer, hyaline, slightly thick-walled, swollen with rounded apex, $35-42 \times 12-18 \mu$.

The voucher specimen has been deposited in the Mycological Herbarium of Burdwan Raj College (BRCMH 7911), Burdwan, West Bengal, India and the duplicate material in the herbarium of the Division of Mycology and Plant Pathology, Indian Agricultural Research Institute, New Delhi, India.

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1. Bilgrami, K. S., Jamaluddin and Rizwi, M. A., *Fungi of India (Part I)*, Today & Tomorrow's Printers and Publishers, New Delhi, 1979.
2. Bilgrami, K. S., Jamaluddin and Rizwi, M. A., *Fungi of India (Part II)*, Today & Tomorrow's Printers and Publishers, New Delhi, 1981.

SAPONINS AND LEUCOANTHOCYANINS IN CASSIA L.

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SAPONINS and leucoanthocyanins have been shown to be of great taxonomic interest in a wide variety of taxa¹⁻⁴. The present article deals with the distribution of saponins and leucoanthocyanins in 21 species of *Cassia* L, and its systematic significance. Saponins were earlier detected in *C. auriculata*, *C. fistula*, *C. italica*, *C. occidentalis*, *C. sophera* and *C. nodosa*⁵ and leucoanthocyanins only in *C. roxburghii*⁶. These species, except *C. italica*, have been reinvestigated here in view of the conflicting reports on the occurrence of saponins^{5, 8}.

The materials used in the present study (table 1) were collected in Bangalore. Voucher specimens are in the Herbarium of the authors' department.

Saponin Test A and Leucoanthocyanin Test A of Gibbs⁷ were used to detect the respective compounds.

The distribution of saponins and leucoanthocyanins

in different parts of the species studied is given in table 1.

The presence of saponins in the species reported to be positive earlier⁵ is confirmed in the present study. There was no correlation between flower colour or habit and the presence of saponins. Since saponins were found in either the leaves or seeds in some species and in both in others, it is necessary to study different parts of the plant to determine their taxonomic distribution. In the present study, quantitative differences from sample to sample were noticed for most species. *C. auriculata* and *C. sophera* were reported⁸ to be saponin-negative while these species were found to be saponin positive⁵ as in the present study. This indicates that interpopulation variation in the presence of saponins does occur. The kind of variation found in the presence of saponins in *Cassia* is not uncommon for secondary compounds in plants⁹. Because of such variation, it is necessary to study several populations before the genetic potentialities for the synthesis of a

Table 1 Distribution of saponins and leucoanthocyanins in *Cassia* species

Species	Saponins		Leucoanthocyanins	
	Leaves	Seeds	Leaves	Flowers
1. <i>C. kleinii</i> W. & A.	-	+	-	-
2. <i>C. mimosoides</i> L.	-	+	-	-
3. <i>C. hirsuta</i> L.	+	+	-	-
4. <i>C. sophera</i> L.	+*	+	-	-
5. <i>C. tora</i> L.	+	-	-	-
6. <i>C. auriculata</i> L.	+*	+	-	-
7. <i>C. montana</i> Heyne ex Roth	-	-	-	-
8. <i>C. alata</i> L.	-	-	-	-
9. <i>C. occidentalis</i> L.	+*	+	-	-
10. <i>C. bicapsularis</i> L.	+	+	-	-
11. <i>C. fruticosa</i> Mill. (= <i>C. bacillaris</i> L.)	+	+	-	-
12. <i>C. carnaul</i> Spreng.	-	-	-	-
13. <i>C. fistula</i> L.	+*	+	-	-
14. <i>C. timorensis</i> DC.	-	-	-	-
15. <i>C. spectabilis</i> DC.	-	+	-	-
16. <i>C. siamea</i> Lam.	+	+	-	-
17. <i>C. nodosa</i> Buch.- Ham ex Roxb.	+*	+	+	+
18. <i>C. javanica</i> L.	+	+	+	+
19. <i>C. grandis</i> L.f.	+	+	+	+
20. <i>C. renigera</i> Wall. ex benth.	+	+	+	+
21. <i>C. roxburghii</i> DC. (= <i>C. marginata</i> Roxb.)	+	+	+*	+*

+ : present; - : absent; * : reinvestigated

particular compound are inferred in a species since genetic potentialities alone should be considered as of taxonomic importance and not the variation which is often inconsistent⁹.

Leucoanthocyanins were absent from the herbaceous and semi-woody species of *Cassia* studied here. This observation is in conformity with the earlier findings on other leguminous taxa^{1,2}. Leucoanthocyanins were absent from the yellow-flowered species of *Cassia*, these are known to contain a large number of anthraquinones, xanthone, cassiixanthone⁴ or barakol¹⁰. The importance of leucoanthocyanins in the systematics of the genus is emphasized by their correlation with woodiness and pink to red flowers. The taxonomic importance of a character increases with its correlatability with other characters¹¹. In the Fabaceae, the absence of leucoanthocyanins was also correlated with epulvinate condition and with affliction by *Uromyces*¹². In the Caesalpinaceae such information is yet unavailable.

C. montana, *C. alata*, *C. carnavall* and *C. timorensis* did not contain either saponins or leucoanthocyanins while *C. nodosa*, *C. javanica*, *C. grandis*, *C. renigera* and *C. roxburghii* were positive for both the compounds. Nevertheless, the rest of the data indicate that the occurrence of saponins and leucoanthocyanins in *Cassia* is independent of each other.

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1. Bate-Smith, E. C., *Proc. Linn. Soc.*, 1958, **169**, 198.
2. Bate-Smith, E. C., *J. Linn. Soc., (Bot.)*, 1962, **58**, 95.
3. Harborne, J. B., In: *Chemotaxonomy of the Leguminosae*. (eds) J. B. Harborne, D. Boulter and B. L. Turner, Academic Press, New York, 1971, p. 257.
4. Harborne, J. B., In: *Chemotaxonomy of the Leguminosae*, (eds) J. B. Harborne, D. Boulter, and B. L. Turner, Academic Press, New York, 1971, p. 257.
5. Rizvi, S. A. I., Lal, J. and Gupta, P. C., *Phytochemistry*, 1971, **10**, 670.
6. Adinarayana, D. and Seshadri, T. R., *Indian J. Chem.*, 1966, **4**, 73.
7. Gibbs, R. D., *Chemotaxonomy of flowering plants*. McGill-Queen's University Press, Montreal, 1974, vol. I, pp. 70, 78.
8. Basalingappa, H. L. and Pathak, C. H., *USDA Forest Research Paper NE 201*, Upper Derby, PA., 1971.

9. Rao, C. K., *Curr. Sci.*, 1983, **52**, 824.
10. Hassanali, A., King, T. J. and Wallwork, S. C., *J. Chem. Soc.*, 1969, **12**, 678.
11. Cronquist, A., *The evolution and classification of flowering plants*. Thomas Nelson, London, 1968, p. 3.
12. El-Gazzar, A., In: *Advances in legume systematics*. (eds) R. M. Polhill and P. H. Raven, Royal Botanic Gardens, Kew, 1981, p. 979.

STUDIES IN NEMATOPHAGOUS FUNGI. XI: *MERISTACRUM ASTEROSPERMUM*—A NEW RECORD FROM INDIA

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WHILE studying nematophagous fungi of Varanasi, a member of Zygomycetes was encountered attacking nematodes. A species of *Meristacrum* is reported for the first time from India as endoparasite of nematode. Only two species of *Meristacrum* are known to parasitize nematodes. *M. asterospermum* has been reported by Drechsler¹ and redescribed by Davidson and Barron². McCulloch³ has described *M. pendulatum* from soils of Queensland. The characters which differentiate the two species are the division of the conidiophore into a number of cells and the shape of conidia. In *M. asterospermum*, only the terminal portion of the conidiophore is divided into cells whereas in *M. pendulatum*, the whole length of the conidiophore is divided into cells.

A detailed account of the chief structural features of *M. asterospermum* Drechsler has been given in this paper. The fungus was isolated using baited plates technique described by Barron⁴. Petri plates containing 2% maize meal agar were baited with heavy suspension of nematodes. Then soil samples were sprinkled on plates. After incubating for a week or so at 25 ± 2°C, the plates were examined. A large number of nematodes were found infected with the fungus.

Meristacrum asterospermum Drechsler: Primary infec-