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INFRARED SPECTROSCOPIC STUDIES IN SOME CLEOME SPECIES

VISHAL T. APARADH* AND B. A. KARADGE

IR spectroscopic study was conducted on the leaves of five species of *Cleome* (viz. *Cleome viscosa* L., *C. chelidonii* L.f., *C. gynandra* L., *C. speciosa* Raf and *C. simplicifolia* (Camb.) Hook f. & Thoms). All studied species showed similar nature of absorption peak at different transmission percentage. Most of the functional groups observed as per their peaks are similar in all five species with little bit difference in wavenumbers. CH₂ Stretching, C=O Stretching, N-H Bending, PO₂- Stretching, C-O Stretching, C-H Stretching, OH and C-H Stretchings are the various functional groups observed. Protein, lipid, amino acid and nucleic acids composition of five *Cleome* species investigated appears to be almost alike which has been evidenced from the present IR spectroscopic studies. This can be taken as one of the parameter for chemotaxonomic support of position of species in the genus *Cleome*.

KEYWORDS: IR spectroscopic study, Cleome species, India

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AUTHOR AFFILIATIONS: Email : Department of Botany, Shivaji University, Kolhapur. (M.S.) India. 416 004. *vishu1415@gmail.com

1. INTRODUCTION

Present investigation have carried out for studying relationship among selective species of Cleome viz. Cleome chelidonii L.f., C. speciosa Raf., C. gynandra L., C. simplicifolia (Camb.) Hook f. & Thoms and C. viscosa L. All these species are herbs growing at same locality but in different soil types as Cleome chelidonii grows vigorously in moist places and also in the rocky regions, while C. simplicifolia and C. viscosa grow luxuriantly in the black soil in rainy season. Cleome simplisifolia has very short life cycle up to 3-4 months only. C. viscosa and C. gynandra grow throughout the year but more vigorously during rainy season. C. gynandra grows predominantly in waste places along waste water. C. speciosa is cultivated species growing widely in shadow places in the red soil particularly during rainy season. It is famous for its beautiful showy inflorescence and hence cultivated in gardens.

2. MATERIALS AND METHODOLOGY

The spectrum of a dried leaf powder of *Cleome* species as sample was determined by an alkali halide pellet method [1]. About 1-3 mg of substance and same amount of KBr were homogenized in mortar with pestle. Dried this mixture for removing moisture and then pressed under high pressure (4-5 ton) at room temperature into a small disc. Pellets were formed. It was kept under IR Spectroscopy to get entire spectrum. As KBr does not absorb infrared radiation in the region 400-4000 cm⁻¹ a complete spectrum of dried powder was obtained.

3. RESULTS

IR spectroscopic study was conducted on the leaves of five species of *Cleome* (Table 1 and Fig 1) (*Cleome viscosa* L., *C. chelidonii* L.f., *C. gynandra* L., *C. speciosa* Raf and *C. simplicifolia* (Camb.) Hook f. & Thoms). The data obtained was analyzed according to Kaur [2].

Cleome viscosa and *C. simplicifolia* have shown protein component having functional groups N-H stretching and O=C-N bending. Instead of O=C-N bending, there is a protein component having C=O bending functional group in *C. chelodonii*. However, in *C. speciosa* leaf protein component has many functional groups (N-H stretching, C=O stretching, C-N stretching, N-H bending and C=O bending) except O=C-N bending. This criterion may be helpful for interspecies differentiation.

In IR spectroscopy, the lipids samples show many different functional groups such as asymmetrical CH_2 stretching, asymmetrical $(CH_3)_3N$ + bending, CH_2 wagging band progression, asymmetrical CO-O-C Stretching and asymmetrical PO₂- stretching. In all *Cleome* species, asymmetrical CH_2 stretching, asymmetrical $(CH_3)_3N$ + bending and CH_2 wagging band progression are the

common functional groups of lipids. However, lipids having asymmetrical CO-O-C stretching and asymmetrical PO₂stretching as functional groups appear to be helpful in taxonomy of species. In *Cleome viscosa* and *C. speciosa* these two functional groups of lipids are absent. While in *C. gynandra* there is presence of lipids containing CO-O-C stretching functional group. Lipids having asymmetrical PO₂- stretching as a functional group appear to be present in *C. chelidonii* and *C. simplicifolia*.

IR spectroscopy of biological sample shows both essential and nonessential type of amino acids having many different functional groups. The IR spectroscopy of *Cleome* species showed the presence of Serine having O-H Bending functional group, Lysine with NH₃+ and NH₃+ bending type and Glutamic acid having symmetric CO₂- stretching as functional groups. Lysine with NH₃+ bending and glutamic acid with symmetric CO₂- are common functional amino acids observed in all *Cleome* species. Serine with O-H Bending functional group is present in *C. viscosa, C. gynandra* and *C. speciosa* species. In *Cleome gynandra* there is presence of only lysine with NH3+functional group. However, there is absence of above two uncommon functional groups in *Cleome chelidonii* and *Cleome simplicifolia*.

Nucleic acids, DNA (deoxyribonucleic acid) and RNA [ribonucleic acid] and proteins make up the most important macromolecules. Nucleic acids are now known to be found in all life forms including bacteria, archaea, mitochondria, chloroplasts, viruses and viroids. All living cells and organelles contain both DNA and RNA, while viruses contain either DNA or RNA, but usually not both (Brock and Madigan, 2009). The basic component of biological nucleic acids is the nucleotide, each of which contains a pentose sugar (ribose or deoxyribose), a phosphate group, and a nucleobase. All these are having CH₂ stretching, C=O stretching, N-H bending, symmetric and asymmetric RNA PO₂- stretching, different RNA ribose C-O stretching with different wavenumbers and DNA ribose C-O stretching as functional groups. In *Cleome* species, CH₂ stretching, C=O stretching and N-H bending, functional groups of nucleic acids are common. Asymmetric RNA PO₂- stretching functional group of nucleic acid is found only in Cleome chelidonii and Cleome simplicifolia. However nucleic acid containing symmetric RNA PO₂- functional group is observed only in Cleome speciosa. In case of nucleic acid with RNA ribose C-O stretching functional groups two different wavenumbers are observed (1015 and 1038), it is 1016.14 in *C. viscosa*, 1035.06 in *C. chelidonii* and 1033.92 in C. simplicifolia with different wavenumber. In C. viscosa there is also presence of DNA ribose C-O stretching at 1016.14 wavenumber. It is absent in other species.

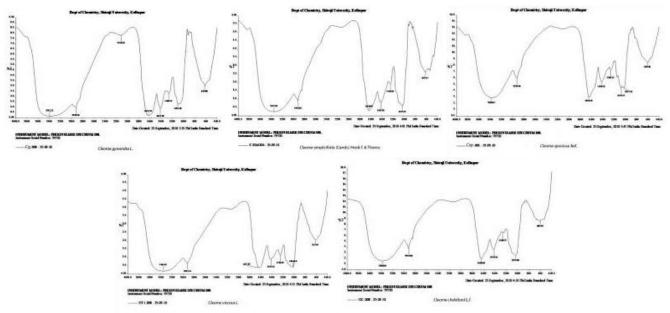


Fig. 1 Comparative infrared spectrum studies in some cleome species

	Functional group		Std wavenumber range cm ⁻¹	Wavenumber as per species (cm ⁻¹)				
Compounds				Cleome viscosa L.	Cleome chelidonii L.f.	Cleome gynandra L.		Cleome simplicifolia (Camb.) Hook f. & Thoms
Proteins	N-H Stretching		3300	3361.02	3380.96	3391.21	3380.62	3353.93
	C=0 Stretching		1653	-	-	-	1650.79	-
	C-N Stretching		1653	-	-	-	1650.79	-
	N-H Bending		1653	-	-	-	1650.79	-
	O=C-N Bending		627	615.43	-	619.08	-	622.92
	C=O Bendi	ng	600	-	603.02	-	605.06	-
Lipids	CH ₂ Stretcl	hing as	2920	2922.71	2914.96	2926.58	2926.58	2918.83
	(CH3)3N+	Bending as	1405	1419.16	1419.16	1415.28	1424.21	1423.03
	CH ₂ Waggi	ng Band Progression	1400-1200	1264.15	1248.65	1260.27	1264.15	1248.65
	CO-O-C Str	etching as	1170	-	-	1101.50	-	-
	PO ₂ -Stretc	hing as	1228	-	1248.65	-	-	1248.65
Amino acids	Serine	O-H Bending	1350-1250	1264.15	-	1260.27	1264.15	-
	Lysine	NH ₃ +	1100	-	-	1101.50	-	-
	Glutamic acid	CO2 ⁻ Stretching s	1415	1419.16	1419.16	1415.28	1424.21	1423.03
	Lysine	NH3 ⁺ Bending	1645-1610	1637.03	1640.04	1627.75	1650.79	1638.09
Nucleic acids	CH ₂ Stretching		2960-2850	2922.71	2914.96	2926.58	2926.58	2918.83
	C=O Stretching		1660-1655	1637.03	1640.04	1627.75	1650.79	1638.09
	N-H Bending		1660-1655	1637.03	1640.04	1627.75	1650.79	1638.09
	RNA PO ₂ - Stretching as		1244	-	1248.65	-	-	1248.65
	RNA P0 ₂ - Stretching s		1084	-	-	-	1074.27	-
	RNA ribose C-O Stretching		1038	-	1035.06	-	-	1033.92
	RNA ribose C-O Stretching		1015	1016.14	-	-	-	-
	DNA ribose C-O Stretching		1015	1016.14	-	-	-	-
Other	C-H Stretching		3350	3361.02	3380.96	3391.21	3380.62	3353.93
	ОН		3350	3361.02	3380.96	3391.21	3380.62	3353.93
	C-H Stretching		2100	-	-	2116.68	-	-

as = asymmetric; s = symmetric

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4. CONCLUSIONS

It was found that all the species studied showed similar nature of absorption peaks but at different transmission percentage. Most of the functional groups observed as per their peaks are similar in all five species with little bit difference in wavenumbers. CH₂ stretching, C=O stretching, N-H bending, PO₂⁻ stretching, C-O stretching, C-H stretching, OH and C-H stretching are the various functional groups observed. Protein, lipid, amino acid and nucleic acids composition of five *Cleome* species investigated appears to be almost alike which has been evidenced from the present IR spectroscopic studies. This can be taken as one of the parameters for chemotaxonomic support of position of species in the genus *Cleome*.

5. References

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