

'The GC MS Analysis of Ethyl Acetate Extract of One Herbal Plant, *Cleome gynandra* ('Gynandropsispentaphylla')

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Abstract

The present study deals with the GC MS analysis of one medicinal plant, '*Cleome gynandra*'. This plant has many ethno-medicinal uses such as immune-modulator, antioxidant, anti-carcinogenic, analgesic, worm infection, ear infections. The plant *Cleome gynandra* was collected from the nearby paddy fields at Chengalpattu, Tamil Nadu. The ethyl acetate extract of the aerial parts of the plant was subjected to GC MS study following standard protocols. It was observed that some very important molecules such as n-Hexadecanoic acid, 4-(2,4-Dimethylcyclohex-3-enyl)but-3-en-2-one, 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, 9,12,15-Octadecatrienoic acid, methyl ester, (Z,Z,Z)-, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Methyl stearidonate, dl-.alpha.-Tocopherol, Campesterol, Stigmasterol, .beta.-Sitosterol, i-Propyl 9,12,15-octadecatrienoate, which have important medicinal roles. The presence of these molecules helps the plant in curing the diseases that are claimed by ethno-medicine. Further work is needed to establish the molecular mechanism of action of these metabolites.

Keywords: GC MS, Ethyl acetate, n-Hexadecanoic acid, 4-(2,4-Dimethylcyclohex-3-enyl)but-3-en-2-one, 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, 9,12,15-Octadecatrienoic acid, methyl ester, (Z,Z,Z)-, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Methyl stearidonate, dl-.alpha.-Tocopherol, Campesterol, Stigmasterol, .beta.-Sitosterol, i-Propyl 9,12,15-octadecatrienoate.

INTRODUCTION

Gynandropsispentaphylla is post rainy season weed with characteristic white flowers, known as Cat's whiskers. This plant has many medicinal roles, such as immune-modulator, antioxidant, anti-carcinogenic, analgesic, worm infection, ear infections. Mishra *et al*, 2011 and Adhikari *et al*, 2018; Adhikari *et al*, 2014, have reviewed the various pharmacological roles of this plant. The antinociceptive and anti-inflammatory activities of leaves of this plant were reported by Mule *et al*, 2008. The anticancer activity of this plant on Ehrlich's ascites carcinoma was studied by Bala *et al*, 2010. Saravanan *et al*, 2017 have studied the GC MS pattern, cytotoxic and antioxidant roles of ethyl acetate extract of this plant. The immune-modulatory role of *Cleome gynandra* was reported by Koriet *et al*, 2009. The present work reports the GC MS pattern of the ethyl acetate extracts of *Cleome gynandra* whole plant. This is in continuation of our endeavour to establish the medicinal efficacy of the herbal and traditional systems of Ayurveda, Sidhha and Unani of medicine (Priyadarshini *et al*, 2017; Jayakumari *et al*, 2017; Rao *et al*, 2018; Vijayalakshmi and Rao, 2019; Yuvaraj *et al*, 2019;

Mutteviet *al*, 2019, Rao *et al*, 2019; Mutteviet *al*, 2020; Vijayalakshmi and Rao, 2020; Janakiet *al*, 2021).

MATERIALS AND METHODS

The plant *Cleome gynandra* was collected from the nearby paddy fields at Chengalpattu, Tamil Nadu. The plant was identified by a qualified botanist at Chennai. The ethyl acetate extract of the shade dried whole plant was collected after 48 h of soaking. The extract was evaporated and the dried powder was used for GC-MS analysis by standard procedures.

GC-MS Procedure

Instrument: GC (Agilent: GC: (G3440A) 7890A. MS/MS: 7000 Triple Quad GCMS) was equipped with MS detector.

Sample Preparation

About 100 ml sample was dissolved in 1 ml of suitable solvents. The solution was stirred vigorously using vortex stirrer for 10 s. The clear extract was determined using GC for analysis.

GC-MS Protocol

Column DB5 MS (30 mm × 0.25 mm ID × 0.25 μm, composed of 5% phenyl 95% methylpolysiloxane), electron impact mode at 70 eV; helium (99.999%) was used as carrier gas at a constant flow of 1 ml/min injector temperature 280°C; auxiliary temperature: 290°C ion-source temperature 280°C.

The oven temperature was programmed from 50°C (isothermal for 1.0 min), with an increase of 40°C/min, to 170°C (isothermal for 4.0 min), then 10°C/min to 310°C (isothermal for 10 min) fragments from 45 to 450 Da. Total GC running time is 32.02 min. The compounds are identified by GC-MS Library (NIST and WILEY).

RESULTS AND DISCUSSION

The results of the GC-MS analysis of the whole plant ethyl acetate extract, along with the possible medicinal role of each molecule of *Cleome gynandra* extract are tabulated in Table 1. Figure 1 represents the GC-MS profile of ethyl acetate extract of the whole plant of *Cleome gynandra*. The identification of metabolites was accomplished by comparison of retention time and fragmentation pattern with mass spectra in the NIST spectral library stored in the computer software (version 1.10 beta, Shimadzu) of the GC-MS along with the possible pharmaceutical roles of each bio molecule as per Dr. Duke's Phytochemical and ethnobotanical data base (National Agriculture Library, USA) and others as shown in Table 1. The results indicates in Table 1 clearly indicate the presence of some important biomolecules, such as n-Hexadecanoic acid, 4-(2,4-Dimethylcyclohex-3-enyl)but-3-en-2-one, 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, 9,12,15-Octadecatrienoic acid, methyl ester, (Z,Z,Z)-, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Methyl stearidonate, dl-α-Tocopherol, Campesterol, Stigmasterol, .beta.-Sitosterol, i-Propyl 9,12,15-octadecatrienoate. These molecules have far reaching medicinal role which corroborate well with the medicinal properties of *Cleome gynandra*.

CONCLUSION

The GC MS profile indicated the presence of some important biomolecules which could contribute to the medicinal role of *Cleome gynandra*.

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REFERENCES

1. Mishra, S. S., Moharana, S. K., Dash, M.R. (2011) Review on *Cleome gynandra*. *Int J Res Pharm Chem*, 1(3), 681-689.
2. Adhikari, P. P., Paul, S. B. (2018) Medicinally important plant *Cleome gynandra*: a phytochemical and pharmacological explanation. *Asian J Pharm Clin Res*, 11(1), 21-29
3. Adhikari, P. P., Paul, S. B., DuttaChoudhury, M., Choudhury, S. (2014) GC-MS Studies on the Steam-distillate of the Medicinally Important Plant *Cleome gynandra* L. *International Journal of Applied Research and Studies*, 3(8), 1-5
4. Mule, S. N., Ghadge, R. V., Chopade, A. R., Bagul, B. A., Patil, S. B., Naikwade, N. S. (2008) Evaluation of anti-nociceptive and anti-inflammatory activity of leaves of *Gynandropsis pentaphylla*. *J Herbal Med Toxicol*, 2, 41-4.
5. Bala, A., Kar, B., Haldar, P. K., Mazumder, U. K., Bera, S. (2010) Evaluation of anticancer activity of *Cleome gynandra* on ehrlich's ascites carcinoma treated mice. *J Ethnopharmacol*, 129, 131-134.
6. Saravanan, R., Pemaiah, B., Narayanan, M., Ramalingam, S. (2017) Gas chromatography-mass spectrometry analysis, *in vitro* cytotoxic and antioxidant efficacy studies on *Cleome gynandra* L. (leaves): A traditional drug source. *Asian J Pharm Clin Res*, 10(10), 84-89
7. Kori, M. L., Gaur, K., Dixit, V. K. (2009) Investigation of immunomodulatory potential of *Cleome gynandra* Linn. *A. J of Pharmaceutical and Clinical Research*, 2(1), 35-39.
8. GomathiPriyadarshini, Arul Amutha Elizabeth, Jacintha Anthony, Mudiganti Ram Krishna Rao, Prabhu. K., Aiswarya Ramesh, Vani Krishna. (2017) The GC MS analysis of one medicinal plant, *Premnatomentosa*. *Journal of Pharmaceutical Sciences and Research*, 9(9), 1595-1597
9. Jayakumari, S., Prabhu, K., Mudiganti Ram Krishna Rao, Bhupesh, G., Kumaran, D., Aishwariya Ramesh. (2017) The GC MS Analysis of a Rare Medicinal Plant *Aloe barbadensis*. *J. Pharm. Sci. & Res.* 9(7), 1035-1037
10. Rao, M. R. K., Vijayalakshmi, N. (2018) Preliminary phytochemical and GC MS analysis of different extracts of *Sphaeranthus indicus* leaves. *Indo American J of Pharmaceutical Sciences*, 5(3), 1511-1520
11. Vijayalakshmi, N., Mudiganti Ram Krishna Rao. (2019) The antioxidant studies of two medicinal plants, *Sphaeranthus indicus* and *Psophocarpus tetragonolobus*. *Asian J of pharmaceutical and Clinical Res*, 12(1), 321-327.
12. Yuvaraj, R., Mudiganti Ram Krishna Rao, Prabhu, K., Lakshmisundram, R., SampadShil, Sathish Kumar, M., Vijayalakshmi, N. (2019) The GC MS study of one medicinal plant, *Stachytarpheta indica*. *Drug Invention Today*, 12(9), 1665-1669
13. Muttevi Hyagreva Kumar, Prabhu, K., Mudiganti Ram Krishna Rao, Lakshmisundram, R., SampadShil, Sathish Kumar, M., Vijayalakshmi, N. (2019) The GC MS study of one medicinal plant, *Dodonaea angustifolia*. *Drug Invention Today*, 12(9), 1661-1664
14. Mudiganti Ram Krishna Rao, Vijayalakshmi, N., Prabhu, K., Sathish Kumar, M. (2019) The gas chromatography-mass spectrometry study of *Moringa oleifera* seeds. *DIT*, 12(10), 2172-2175

15. MutteviHyagreva Kumar, Prabhu, K., Mudiganti Ram Krishna Rao, Lakshmisundram, R., SampadShil, Sathish Kumar, M., Vijayalakshmi, N.(2020) The GC MS study of one medicinal plant, *AristolochiaIndica* .DIT, 12(12), 2919-2923.

16. Vijayalakshmi, N., Mudiganti Ram Krishna Rao. (2020) ‘Preliminary phytochemical and antioxidant studies of leaf extracts of one medicinal plant, *Vitexnegundo*’ .RJPT, 13(5), 2167-2173

17. Janaki C. S.; Prabhu K., Mudiganti Ram Krishna Rao, VenkatRamaiah, ShrutiDinkar, Vijayalakshmi, N., Kalaivannan. J. (2021) The GC MS analysis of Ethyl acetate extract of *Merremiaemerginata*’.Ind J of Natural Sciences, 12(67), 33638-33646

18. Dr.Duke’s Phytochemical and Ehnobotanical Databases.U.S. Department of Agriculture, Agricultural Research Service.1992-2016. Dr. Duke’s Phytochemical and Ethnobotanical Databases. Home Page, <http://phytochem.nal.usda.gov/> <http://dx.doi.org/10.15482/USDA.ADC/1239279>

Qualitative Compound Report

Data File	280121036.D	Sample Name	Gynandropsis pentaphylla
Sample Type		Position	123
Acq Method	GC Screening New Method.M	Acquired Time	31-01-2021 AM01:22:35
Comment			

User Chromatogram

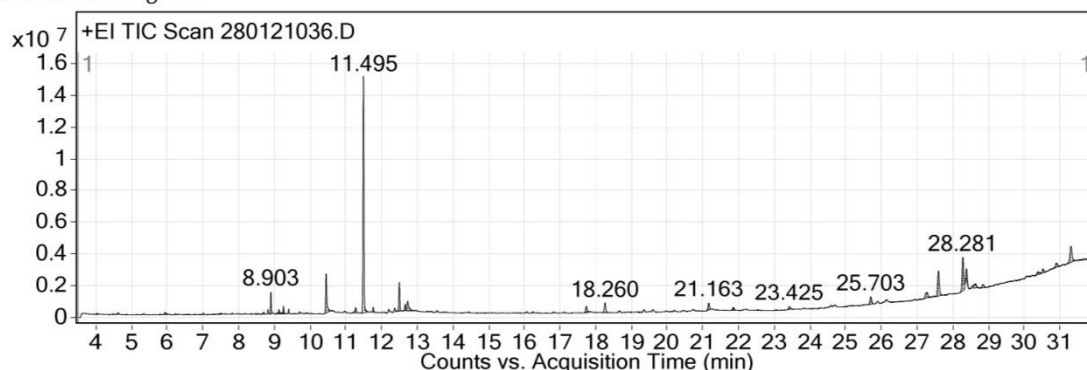


Figure 1. Shows the GC MS profile graph of ethyl acetate extract of *Cleome gynadra*

Table1. Indicates the retentions time, types of possible compound, molecular formula, molecular mass, percentage peak area and the possible medicinal roles of each compound as shown in the GC MS profile of *Cleome gynadra*

Ret. Time	Compound	Mol. Formula	Mol. Mass	% Peak area	Possible Medicinal role
8.90	Bicyclo[3.1.1]heptane, trimethyl-	2,6,6-C ₁₀ H ₁₈	138.1	2.83	Not Known
10.45	n-Hexadecanoic acid	C ₁₆ H ₃₂ O ₂	256.2	7.40	Acidifier, Arachidonic acid Inhibitor, Increases

					Aromatic Amino acid decarboxylase activity, Inhibits production of uric acid, Urine acidifier, Anaphylactic, Arylamine N acetyltransferase inhibitor, decreases norepinephrine production, Down regulates nuclear and cytosol androgen reuptake, GABA-nergic, Increase NK cell activity, inhibits production of tumor necrosis factor, Myo-neuro-stimulator
11.28	4-(2,4-Dimethylcyclohex-3-enyl)but-3-en-2-one	C ₁₂ H ₁₈ O	178.1	1.11	Decreases endothelial Leukocyte adhesion, Decreases endothelial Platelet adhesion, endocrine tonic, endothelium derived relaxing factor promoter, enter relaxant, enteromotility enhancer
11.50	Cyclohexanol, 5-methyl-2-(1-methylethyl)-, (1.alpha.,2.beta.,5.alpha.)-(./-.)-	C ₁₀ H ₂₀ O	156.2	37.67	Not Known
12.66	Ethyl 9,12,15-octadecatrienoate	C ₂₀ H ₃₄ O ₂	306.3	1.14	Not known
12.73	9,12,15-Octadecatrienoic acid, methyl ester, (Z,Z,Z)-	C ₁₉ H ₃₂ O ₂	292.2	2.81	Acidifier, Arachidonic acid Inhibitor, Increases Aromatic Amino acid decarboxylase activity, Inhibits production of uric acid, Urine acidifier,
17.74	Methyl 16-hydroxy-hexadecanoate	C ₁₇ H ₃₄ O ₃	286.3	1.61	Catechol-O-Methyl-Transfearse inhibitor, Increases Glutathione-s-Transferase Activity, Decrease Glutamate Oxaloacetate transaminase activity, Decreases Glutamate pyruvate transaminase, Glycosyltransferase inhibitor, Glutathione-S-Transfearse inhibitor, Increases glyoxalate

					transamination, Reverse transcriptase inhibitor,
18.26	2-((Octan-2-yloxy)carbonyl)benzoic acid	C16H22O4	278.2	2.38	Acidifier, Arachidonic acid Inhibitor, Increases Aromatic Amino acid decarboxylase activity, Inhibits production of uric acid, Urine acidifier,
21.16	Methyl stearidonate	C19H30O2	290.2	1.91	Catechol-O-Methyl-Transfearse inhibitor, Methyl donor
25.70	dl-.alpha.-Tocopherol	C29H50O2	430.4	2.08	Tocopherol synergist, 5 alpha reductase inhibitor, Alpha agonist, Alpha amylase inhibitor, Alpha glucosidase inhibitor, HIF-1 alpha inhibitor, Ikappa B-alpha phosphorylation inhibitor, Increase alpha mannosidase activity, Interleukin 1-alpha inhibitor, Testosterone-5-Alpha-Reductase-Inhibitor, TNF- alpha inhibitor
27.28	Campesterol	C28H48O	400.4	2.43	Plant steroid use as food additive and has cholesterol lowering role
27.60	Stigmasterol	C29H48O	412.4	7.49	Precursor of progesterone act as intermediate in the biosynthesis of androgens and estrogens Antiosteoarthritic, antihypercholestrolemic, cytotoxic, antitumor, hypoglycaemic, antimutagenic, antioxidant, anti-inflammatory, Analgesic.
28.28	.beta.-Sitosterol	C29H50O	414.4	9.48	17 beta dehydrogenase inhibitor, androgen blocker, anti-amyloid beta, anticancer, Anti TGF beta, Beta 2- receptor, beta blocker, beta-galactosidase inhibitor, beta-glucuronidase inhibitor
28.38	3,7,11,15-Tetramethyl-2-hexadecen-1-ol	C20H40O	296.3	7.26	Catechol-O-Methyl-Transfearse inhibitor, Methyl donar
28.6	Phytonadione	C31H46	450.	1.53	Not Known

3		O2	4		
28.8 5	Cholest-5-en-3-ol, 24-propylidene-, (3.beta.)-	C30H50 O	426. 4	1.01	Not Known
31.3 1	i-Propyl 9,12,15-octadecatrienoate	C21H36 O2	320. 3	5.69	11Beta HSD inhibitor, 17-beta-hydroxysteroid dehydrogenase inhibitor, 5 HETE inhibitor, 5 HT inhibitor, 8 HETE inhibitor, Anti 5-HT, 5-Lipoxygenase-Inhibitor, 8-HETE-Inhibitor, Aldehyde-Oxidase-Inhibitor, Alcohol-Dehydrogenase-Inhibitor, AHH-Inhibitor, Akt-Inhibitor, Adenylate-Cyclase-Inhibitor, AChE-Inhibitor, Acetyl-CoA-Carboxylase-Inhibitor, ACE-Inhibitor, ABeta-Inhibitor