

'The Gc Ms Analysis Of Ethyl Acetate Extract Of One Herbal Plant, 'Spermacoce Hispida L'

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ABSTRACT

The present study deals with the GC MS analysis of one medicinal plant, 'Spermacoce hispida L'.This plant has wide application in ethnomedical practice. It is used to cure stomach ailments, dandruff, boils and swellings. This plant was collected from nearby hills of Chengalpattu, Tamilnadu. 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, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Squalene, .gamma.-Tocopherolm, dl-.alpha.-Tocopherol, Campesterol, Stigmasterol, .beta.-Sitosterol, E,E,Z-1,3,12-Nonadecatriene-5,14-diol. The medicinal roles of these molecules could contribute to the medicinal roles of the plant as claimed by ethno-medicinal practice.

Keywords : GC MS, Spermacoce hispida , Ethyl acetate, n-Hexadecanoic acid, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Squalene, .gamma.-Tocopherol, dl-.alpha.-Tocopherol, Campesterol, Stigmasterol, .beta.-Sitosterol,

INTRODUCTION

The present works deals with the GC MS analysis of the whole plant of Spermacoce hispida. This plant has wide application in ethno-medical practice. It is used to cure stomach ailments and dandruff. The flowers are used to treat boils and swellings. The seed extracts are used to treat internal injuries of nerves and kidney. It purifies blood and improves vigour of the body(Meti et al, 2013). Anupriya et al, 2016 have reported the GC MS studies on different extracts of Spermacoce hispida L.Anantharaman and Krishnamoorthy, 2017 and Dhevi and Elango, 2015 have reported the

GC MS studies of the chloroform extracts of the seeds of Spermacoce hispida L. The leaves of Spermacoce hispida L have been reported to have cancer therapeutic role by Rathi et al, 2011. This work is in continuation of our work to establish the efficacy of the herbal plants, Ayurvedic and Sidhha medicines. (Priyadarshini et al, 2017; Jayakumari et al, 2017; Rao et al, 2018; Vijayalakshmi and Rao, 2019; Yuvaraj et al, 2019; Muttevi et al, 2019, Rao et al, 2019; Muttevi et al, 2020; Vijayalakshmi and Rao, 2020; Janaki et al, 2021)

MATERIALS AND METHODS

The plant Spermacoce hispida was collected from the nearby hills 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 μ m, 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; auxilary 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 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 DISCUSION

The results of the GC-MS analysis of the whole plant ethyl acetate extract, along with the possible medicinal role of each molecule ofSpermacoce hispida extract are tabulated in Table 1. Figure 1 represents the GC-MS profile of ethyl acetate extract of the whole plant of Spermacoce hispida. 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

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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.n-Hexadecanoic acid, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Squalene, .gamma.-Tocopherol, dl-.alpha.-Tocopherol, Campesterol, Stigmasterol, .beta.-Sitosterol, E,E,Z-1,3,12-Nonadecatriene-5,14-diol etc. These molecules are known to have far reaching medicinal roles as mentioned in Table 1 which corroborate well the medicinal roles of this plant.

CONCLUSION

From the above mentioned resultsit is clear that Spermacoce hispidahas some important medicinal roles. Further work in this regard is warranted.

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Figure 1. Shows the GC MS profile graph of ethyl acetate extract of Spermacoce hispida L

Qualitative Compound Report



Table1. Indicates the retentions time, types of possible compound, molecular formula, molecularmass, percentage peak area and the possible medicinal roles of each compound as shown in the GCMS profile of Spermacoce hispida L

Ret.	Compound	Mol.	Mol	%	Possible Medicinal Role
Time		Formula		Peak	
			Mas	Area	
			S		
8.5	1-Pentadecyne	C15H28	208.	0.98	Not known
			2		
8.90	Bicyclo[3.1.1]heptane, 2,6,6-	C10H18	138.	2.21	Not known
	trimethyl-		1		
10.4	n-Hexadecanoic acid	C16H32O	256.	7.32	Acidifier, Arachidonic acid
5		2	2		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
11.4	Cyclohexanol, 5-	C10H20O	156.	13.8	Not known
9	methyl-2-(1-		2	0	
	methylethyl)-,				
	(1.alpha.,2.beta.,				
	5.alpha.)-(.+/)-				
18.2	2-((Octan-2-	C16H22O	278.	7.54	Acidifier, Arachidonic acid
5	yloxy)carbonyl)benzoic acid	4	2		inhibitor, Increases Aromatic
					Amino acid Decarboxylase
					activity
20.1	Squalene	C30H50	410.	19.5	Plant steroid use as food
9			4	8	additive and has cholesterol
					lowering role
24.7	.gammaTocopherol	C28H48O	416.	0.65	Tocopherol synergist,
0		2	4		PPAR-gamma antagonist
25.5	4,5,6,7-Tetrahydro-	C12H15N	221.	5.5	Not known
1	benzo[c]thiophene-1-	os	1	3	
	carboxylic acid allylamide				
25.6	dlalphaTocopherol	C29H50O	430.	2.1	Tocopherol synergist, 5 alpha
9		2	4	9	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
27.2	Campesterol	C28H48O	400.	4.1	Plant steroid use as food
6			4	4	additive and has cholesterol
					lowering role
27.5	Stigmasterol	C29H48O	412.	7.4	Precursor of progesterone,
8			4	0	acts as intermediate in the
					biosynthesis of androgens
					and estrogens, anti-
					osteoarthritic,
					antihypercholesterolemic,
					cytotoxic, antitumor,
					hypoglycemic,
					antimutagenic, antioxidant,
					anti-inflammatory, analgesic
28.2	.betaSitosterol	C29H50O	414.	8.3	17 beta dehydrogenase
7			4	1	inhibitor, androgen blocker,
					anti-amyloid beta,
					anticancer, Anti TGF beta,
					Beta 2- receptor, beta
					blocker, beta-galactosidase
					inhibitor, beta-glucuronidase
					inhibitor
28.3	3,7,11,15-Tetramethyl-2-	C20H40O	296.	7.2	Oligosaccharide provider
6	hexadecen-1-ol		3	9	
28.6	Phytonadione	C31H46O	450.	0.5	Not known
1		2	4	9	

30.6	E,E,Z-1,3,12-Nonadecatriene-	C19H34O	294.	1.1	anticancer, antidote,
1	5,14-diol	2	3	4	antitumor, Cytochrome-
					P450-2E1-Inhibitor,
					Decreases C-Teleopeptide
					Excretion, Decreases
					Deoxypyridinoline Excretion,
					Decreases Endothilial
					Leukocyte Adhesion,
					Decreases Epinephrine
					Production, Decreases
					Oxalate Excretion