

‘The Gc Ms Analysis Of Ethyl Acetate Extract Of One Herbal Plant, ‘Cyanotis Tuberosa’

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ABSTRACT

The present study deals with the GC MS analysis of one medicinal plant, ‘Cyanotistuberosa’. Not much work was done on this species and it is of interest to learn about the medicinal value since ethnopharmacological reports on this plant are also scanty. The plant was collected from nearby hills of Chengalpattu, Tamilnadu. The ethyl acetate extract of the tuber of the plant was subjected to GC MS study following standard protocols. It was observed that some very important molecules such as 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, n-Hexadecanoic acid, -((Octan-2-yloxy)carbonyl)benzoic acid, Cholesterol, 3H-Isobenzofuran-1-one, 6,7-dimethoxy-3-p-tolylamino- Further probe is warranted to known about the medicinal role of this plant in the light of the present report.

Keywords GC MS, Cyanotistuberosa, 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, n-Hexadecanoic acid,, Cholesterol, 3H-Isobenzofuran-1-one

INTRODUCTION

The plant, Cyanotistuberosa is found in the hill regions and grows during monsoon. The tubers are perennial and they produce fresh aerial shoot every season. No scientific reports are available on the medicinal role of this plant and ethno-botanically also there are very scanty reports on its medicinal

value. Ghosh and Chowdhury, 2016 have reported the pharmacognosy and phytochemical analysis of *Cyanotistuberosa*. Therefore this plant was chosen for our study. The plant along with the tuber was collected from the hills near Chengalpattu, Tamil Nadu, and the ethyl acetate extract of the tuber was subjected to GC MS analysis following standard protocols. This work is in continuation of our endeavor to substantiate the medicinal roles of herbal plants, Ayurvedic and Sidhha medicines. (Priyadarshiniet al, 2017; Jayakumariet al, 2017; Raoet al, 2018; Vijayalakshmi and Rao, 2019; Yuvarajet al, 2019; Mutteviet al, 2019, Raoet al, 2019; Mutteviet al, 2020; Vijayalakshmi and Rao, 2020; Janakiet al, 2021, Perumalet al, 2021).

MATERIALS AND METHODS

The plant, *Cyanotistuberosa* was collected from the nearby hills of Chengalpttu, 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 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 of *Cyanotistuberosa* ethyl acetate extract, along with the possible medicinal role of each molecule are tabulated in Table 1. Figure 1 represents the GC-MS profile of ethyl acetate extract of the whole plant of *Cyanotistuberosa*. 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 ethno-botanical data base (National Agriculture Library, USA) and others as shown in Table 1. Some molecules as represented by the GC MS profile indicated the presence of some important biomolecules such as 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, n-Hexadecanoic acid, -((Octan-2-yloxy)carbonyl)benzoic acid, Cholesterol, 3H-Isobenzofuran-1-one, 6,7-dimethoxy-3-p-tolylamino- Further probe is warranted to known about the medicinal role of this plant in the light of the present report.

CONCLUSION

Thus it can be concluded that due to the presence of these molecules, *Cyanotistuberosa* has some the medicinal roles and it should be explored as a possible medicine. Further work to isolate and understand the molecular mechanism is warranted.

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Qualitative Compound Report

Data File	280121045.D	Sample Name	Cynotis tuberosa
Sample Type		Position	132
Acq Method	GC Screening New Method.M	Acquired Time	31-01-2021 AM06:56:54
Comment			

User Chromatogram

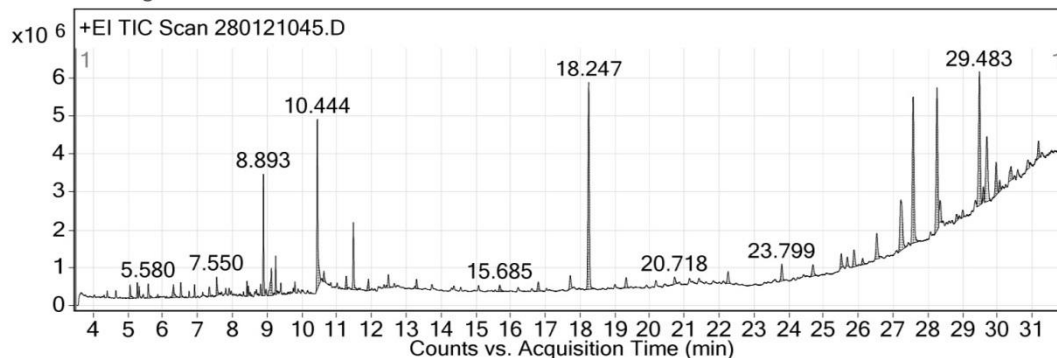


Figure 1. Represents the GC MS graph of ethyl acetate extractCyanotistuberosa

Table 1. 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 Cyanotistuberosa

Ret. Time	Molecules	Mol. Formula	Mol. Mass	% Peak area	Possible Medicinal Role
8.89	1-Hexadecyne	C16H30	222.2	3.91	Not Known
9.12	9-Octadecyne	C18H34	250.3	1.82	Not Known
9.25	3,7,11,15-Tetramethyl-2-hexadecen-1-ol	C20H40O	296.3	1.23	Oligosaccharide provider
10.44	n-Hexadecanoic acid	C16H32O2	256.2	7.98	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
11.48	Cyclohexanol, 5-methyl-2-(1-methylthyl)-, (1.alpha.,2.beta.,5.alpha.)-(+/-)-	C10H20O	156.2	2.93	Not known
18.25	2-((Octan-2-ylloxy)carbonyl)benzoic	C16H22O4	278.2	13.35	Acidifier, Arachidonic acid inhibitor, Increases Aromatic

	acid				Amino acid Decarboxylase activity
23.80	3-Eicosyne	C ₂₀ H ₃₈	278.3	1.16	Not known
25.88	Cholesterol	C ₂₇ H ₄₆ O	386.4	1.33	The precursor for steroid hormone synthesis
26.53	3H-Isobenzofuran-1-one, 6,7-dimethoxy-3-p-tolylamino-	C ₁₇ H ₁₇ NO 4	299.1	2.10	11Beta HSD inhibitor, 17-beta-hydroxysteroid dehydrogenase inhibitor, 5 HETE inhibitor, 5 HT inhibitor, 8 HETE inhibitor, Anti 5-HT, Anti HIV integrase, Aryl hydrocarbon hydroxylase inhibitor, HDL genic, Hematopoietic