ejpmr, 2015,2(4), 219-226

SJIF Impact Factor 2.026



EUROPEAN JOURNAL OF PHARMACEUTICAL AND MEDICAL RESEARCH

<u>www.ejpmr.com</u>

Research Article ISSN 3294-3211

EJPMR

GC-MS ANALYSIS OF ETHYL ACETATE EXTRACT OF SEMECARPUS ANACARDIUM NUTS

*Rajendra Singh and Arun Kakkar

*Govt. Model Science College (Auto.), Dept of Chemistry, South Civil Line Pachpedi, Jabalpur (M.P.) India, 482001.

Article Received on 22/04/2015Article Revised on 13/05/2015Article Accepted on 04/06/2015

*Correspondence for Author Rajendra Singh Govt. Model Science College (Auto.), Dept of Chemistry, South Civil Line Pachpedi, Jabalpur (M.P.) India, 482001

ABSTRACTS

In present investigation phytochemical compounds of *Semecarpus anacardium* nuts using GC-MS technique was carried out to determine the structure of phytochemicals present in it. The mass spectra of the phytocompounds found was matched with the National Institute of Standards and Technology library. Dried SA nut was successively extracted with petroleum ether, ethyl acetate, methanol and water using

soxhlet apparatus. This extracts was purified and stored in refrigerator. The ethyl acetate extract of SA nuts was analyzed by Agilent 7890A GC with 5975MS. Nine compounds were indentified ie benzoic acid, 4-ethoxy ethyl ester, phenol. 2,4- bis (1,1-dimethyl), Eicosane, 7-hexyl. The compounds were identified by comparing their retention time and peak area with literature and structure determination was done by interpretation of mass spectra. Present investigation by GC-MS analysis of ethyl acetate extract of SA is done.

Abbreviation

SA = Semecarpus anacardium GC-MS = Gas Chromatography Mass Spectroscopy TLC = Thin layer Chromatography NIST =National Institute Standard and Technology TDA =Toluene-diamine EI = Electron Ionization PCI = Positive Chemical Ionization NCI = Negative Chemical Ionization RT = Retention Time MF = Molecular Formula MW = Molecular Weight amu = atomic mass unit

KEY WORDS: Bhilwa, Phytochemical, Herbal medicine, Anacardiaceae

INTRODUCTION

Semecarpus anacardium (SA) belongs to Anacardiaceae family commonly known as bhallataka or marking nut.^[1] SA nuts are used in the Ayurveda and Siddha systems of medicine, with various therapeutic properties such as anti-atherogenic effect^[2], antiinflammatory^[3], anti-oxidant^[4], anti-microbial⁵, CNS⁶, hypoglycemic^[7], anticarcinogenic^[8], hyperlipidemic activity.^[9] Reported constituent of SA seeds are blilawanol, anacardoside, in fruits are nicotinic acid, riboflavin, thiamine, arginine, histidine, isoleucine, leucine, lysine, methionene, pheynylanine, threonine, tryptophan, valine, in nuts are tetrahydroamentoflavone, biflanoids A, B, C, 3, 8- binaringenin, 3,8-biliquiritigeninnn, nallaflavanone, oil are anacardiac acid, cardol and catechol.^[10-13] The medicinal properties of this plant were attributed due to its variety of active phytochemical constituent. Although the plant had received a great interest for the phytochemical investigation since many years Various parts of SA were investigated mainly for the presence of major types of phytochemical compounds. Hence the objective of the present study is due to identify the phytochemical constituent of the ethyl acetate extract with the aid of GC-MS technique and to correct the activities with the active constituent present.

MATERIAL AND METHODS

Collection of plant materials

The nuts of SA was collected during the month of September from the tribal area of Ambikapur (Wadrafnagar, Madhna) Chhattisgarh India. The plant was taxonomically identified by Professor Dr. K. P. Sahu, Botony Department, Govt. Model Science College, Jabalpur.

Preparation of extracts Organic solvent of increasing polarity were used extraction according to the methodology of Indian pharmacopoeia.^[14] Dried SA nut was successively extracted with petroleum ether, ethyl acetate, methanol and water using soxhlet apparatus. This extracts was purified and stored in refrigerator.

GC-MS analysis of material

Instruments Agilent 5975C TDA series gas chromatography/mass spectroscopy selective detector system offer high performance and flexibility with many options. Gas chromatograph Agilent 7890 A is the auto sampler, oven temperature is ambient $+4-450^{\circ}$ C and 20/21 negative ramps allowed and mass selective detector includes standard mode- EI, optional mode-PCI, NCI and EI acquisition with CI source, EI Ion source type-non coated inert EI source, Ion source temperature-150°C to 350°C, Quadruple temperature-106-200°C, mass filter-monolithic hyperbolic quadrupole, minimum mass- 1.6µ, maximum mass - 1050µ, mass axis stability-better than 0.10 µ/48h, detector-triple axis.

Method of GC-MS analysis and chromatographic condition Crude ethyl acetate extract (1 μ l) of SA nut was used in GC-MS analysis. GC-MS of ethyl acetate extract was performed using Agilent 7890A. Column used on Agilent (5975C MS) 5% poly siloxane column 30×250 μ m×0.25 μ m size. Oven temperature was programmed as follows: Isothermal temperature was 5°C/min and held for 1.75 min then increased to 275°C at the rate of 8°C/min and kept constant for 5min. The run time was 25min. Ionization of sample components were performed on EI mode (70eV).

Identification of components Interpretation on mass spectrum GC-MS was done using the database of National Institute Standard and Technology (NIST). The spectrum of the NIST library. The name molecular weight and structure of the compounds of the ethyl acetate ascertained.

RESULTS AND DISCUSSION

GC-MS chromatogram of the ethyl acetate extract of SA nuts (Fig-1) showed nine peaks indicating the presence of nine compounds. The chemical compounds identified in the ethyl acetate extract of the nuts of SA are presented in Table-1. GC-MS analysis revealed the presence of benzoic acid, heptadecane 9 hexyl, octadecane 3-ethyl -5 (2-ehtylbutyl), glucobrassicin, eicosane, 7- hexyl. Keeping in view the tremendous pharmacological activities of its constituent, SA is utilized to alleviate the symptoms of variety of diseases. Phenol 2,4 bis (1,1-dimethyl ethyl) has good antibacterial activity.^[15]. The wide spread of availability and extensive literature of SA in India thus makes it an attractive target for further pre-clinical and clinical research. These compounds are repeated first time from GC-MS analysis of SA nuts ethyl acetate extract.

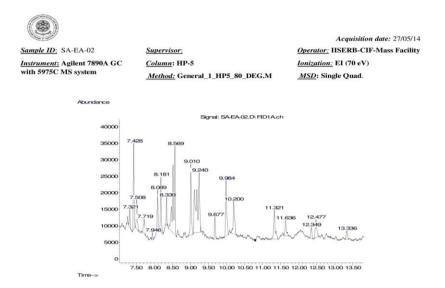


Fig:1. GC of SA nuts crude extract in ethyl acetate

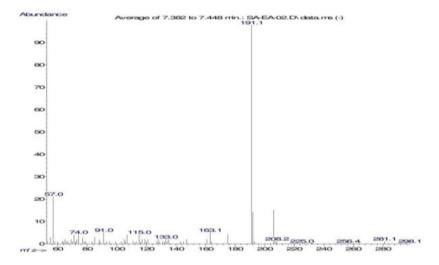


Fig1(a).MS of Phenol, 2,4-bis(1,1-dimethyl ethyl)

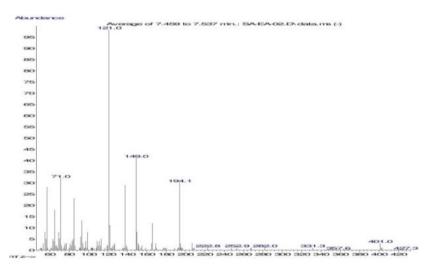
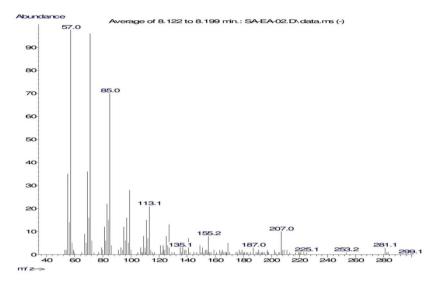
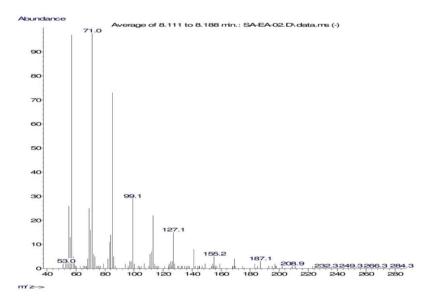


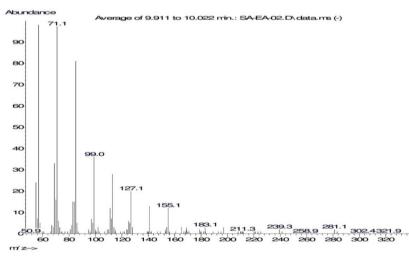
Fig 1(b). MS of Benzoic acid, 4-ethoxy-ethyl ester



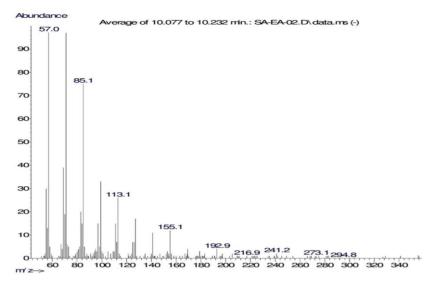














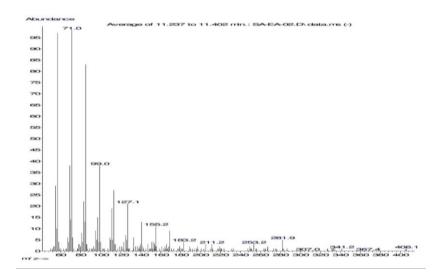


Fig 1 (G). MS of Octadecane, 3-ethyl-5(2-ethyl butyl)

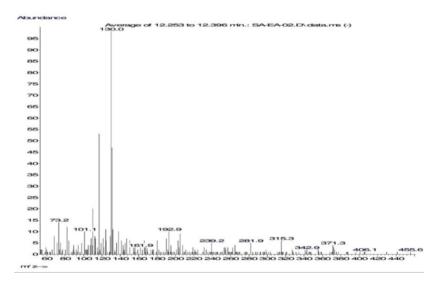


Fig 1 (h). MS of Glucobrassicin

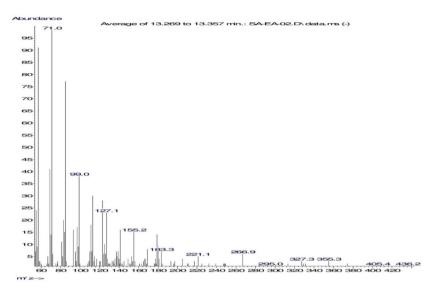


Fig 1 (I). MS of Eicosane, 7-Hexyl

Table 1:	Phytochemicals identified in crude SA nuts extract in ethyl acetate of by GC-
MS	

S. No.	RT	Name of the isolated	Molecular	MW	Peak
	(In min.)	compound	formula	(amu)	% area
1	7.44	Phenol 2,4-bis (1,1-dimethylethyl)	$C_{14}H_{22}O$	206	7.79
2	7.53	Benzoic acid, 4-ethoxy- ethyl ester,	$C_{11}H_{14}O_3$	194	3.60
3	8.14	Icosane	$C_{20}H_{42}$	282	3.51
4	8.19	Eicosane, 2-methyl	$C_{21}H_{44}$	296	5.32
5	9.98	Tetracosane	$C_{24}H_{50}$	338	7.30
6	10.20	Hepatadecane,9-hexyl	C ₂₃ H ₄₈	324	5.19
7	11.32	Octadecane,3-ethyl-5- (2-ethyl butyl)	C ₂₆ H ₅₄	366	4.96
8	12.32	Glucobrassicin	$C_{16}H_{20}N_2O_9S_2$	448	3.06
9	13.51	Eicosane, 7- hexyl	C ₂₆ H ₅₄	366	1.24

ACKNOWLEDGEMEN

The authors wish to thank Principal, Govt. Model Science College and Head of Chemistry Department Govt. Model Science College, Jabalpur (M.P.) for providing the necessary laboratory facilities. Thanks also are due to UGC New Delhi for financial assistance.

REFERENCES

- Khare CP, Semecarpus. In Encyclopedia of Indian medicinal plants, Springer: Germany; 2004.
- Sharma A, Mathur R, Dixit VP. Hypocholesterol activity of nut shell extract of some medicinal plants of Bangladesh, Dhaka Univ. J Pharma Sci 2008; 7:47-52.
- 3. Bhitre MJ, Patil S, Katarial M, Anwikar S, Kadri H. Anti-inflammatory activity of the fruits of *Semecarpus anacardium* Linn. Asian J chem. 2008;20:47-52.

- 4. Sahoo AK, Narayanana N, Sharma S, Rajanb SS, Mukherjee PK, Vitro antioxidant potential of *Semecarpus anacardium* L. Pharmacologyonline 2008;3:27-35.
- Mohanta TK, Patra JK, Rath SK, Pal DK, Thatoi HN. Evalution of antimicrobial activity and Phytochemical screening of oil *Semecarpus anacardium*. Sci Res Essay 2007;2:28-69.
- Farooq SM, Alla TR, Prasad K, Shalam K, Satyanarayna. Semecarpus anacardium stydy on CNS effect of nut milk extract of Semecarpus anacardium. Pharmacogyoline 2007;149-163.
- Arul B, Kothari R, Christine AJ. Hypoglycemic activity of *Semecarpus anacardium* Linn in normal and streptozotocin-induced diabetic rats. Methods find Exp. Clin pharmcol. 2004;7:59-62.
- Veena K, Shanthi P, Sachdanandam P. The biochemical alterations following administration of kalpaamruthaa and *Semecarpus anacardium* in mammary Chem Biol Interact 2006;169:69-78.
- Kaladevi SK, Haseema BHK, Keerthiga G., Shanthi P., Sachdanadam P. Hypolipidemic effect of *Semecarpus anacardium* in high cholesterol fed hyperchlosterolmic rats. Chin J inte Med 2012;3: 2321-2575.
- Murthy, SSN, A biflavone from *Semecarpus anacardium*, Phtochemistry, 1983; 22 (6): 1518-1520
- 11. Murthy, SSN, Confirmation of the structure of jeediflavaone A biflavanone from *Semecarpus anacardium*, phytochemistry, 1984; 23 (4): 925-927.
- 12. Murthy, SSN, Further evidence to the structure of semecarpuflavanone, curr. Science, 1984; 53 : 643-645.
- 13. Murthy, SSN, Semecarpetin, a biflavanone from *Semecarpus anacardium*, Phytochemistry, 1988; 27 (9): 3020-3022.
- Anonymous Pharmacopoeia of India (*The Indian pharmacopoeia*) 2nd Edition, Manager of publication, New Delhi: 1966, PP. 947-948.
- 15. ASH. Abdullah, MES. Mirghani and P. Jamal, Antibacterial activity of Malaysian mango kernel, African Journal of Biotechnology, 2011; 10 (81):18739-18748.