See discussions, stats, and author profiles for this publication at: https://www.researchgate.net/publication/281062356

GC-MS Determination of Bioactive Constituents of Giant African Snail (Archachatina maginata) Haemolymph

	May 2015 //3008-10245964			
CITATIONS		READS		
4		69		
5 author	s, including:			
	Oluwatosin Kudirat Shittu		Prince Ossai	
	Federal University of Technology Minna		Federal University of Technology Minna	
	36 PUBLICATIONS 107 CITATIONS		7 PUBLICATIONS 9 CITATIONS	
	SEE PROFILE		SEE PROFILE	

Some of the authors of this publication are also working on these related projects:



Proteomic studies on the malaria parasite Plasmodium falciparum. View project

GC-MS Determination of Bioactive Constituents of Giant African Snail (*Archachatina maginata*) Haemolymph

*Bashir Lawal, Oluwatosin K. Shittu, Tawakaltu AbdulRasheed-Adeleke, Prince C. Ossai, and Aisha M. Ibrahim

Department of Biochemistry, Federal University of Technology, Tropical Disease Research Unit, PMB 65, Minna, Nigeria

Abstract: Giant African snail (Archachatina maginata) is of high medicinal value, it haemolymph has been used in folk medicine for the treatment of liver disorders, whooping cough, anaemia, constipation, restore vitality and stop bleeding. In tune with this effect, the objective set for the present study is to identify the bioactive constituents of A. marginata haemolymph in order to understand the nature of the principle component responsible for its medicinal property. The haemolymph was extracted from the snail (A. maginata) and subjected to Gas Chromatography-Mass Spectrometry (GC-MS) analysis using a GC-MS (Model: QP2010 PLUS SHIMADZU, JAPAN) comprising a AOC-20i auto-sampler and gas-chromatograph interfaced to a mass spectrometer. GC-MS analysis provided of seven peaks. On comparison of the mass spectra of the constituents with the NIST library twenty six (26) constituents including 7 ester, 7 fatty acid, 5 alcohol, 6 alkane and 1 phthalate were characterized and identified. The presence of various bioactive compounds justifies it uses for various ailments by traditional practitioners. However isolation of individual constituents and subjecting it to biological activity will definitely give fruitful results and helpful to find anew drugs.

Keywords: Snail, Haemolymph, GC-MS analysis, bioactive constituents.

I. Introduction

Natural products, owing to their medicinal value have continued to play a dominant role in the maintenance of human health since ancient times. The search on natural product have led to the discovery of novel drug candidates used against diverse diseases, as over 50% of all modern clinical drugs are of natural product origin [1].

A knowledge of the chemical constituents of natural products is desirable not only for the discovery of therapeutic agents, but also because such information may be of great value in disclosing new sources of bioactive principle for the synthesis of complex chemical substances and for discovering the actual significance of folkloric remedies [2]. Hence a thorough validation of the natural products with medicinal reputation has emerged as a new branch of science emphasizing and prioritizing the standardization of the natural drugs and products because several of the bioactive agents have complementary and overlapping mechanism of action [3]. Several wild animals and their secreation have been used in folklore medicine. The African giant snail (*Archachatina marginata*) is one of the most important minor forest products in West Africa and Nigeria in particular [4].

The haemolymph of A. marginata contains a myriad of metabolites and other constituents which provide a valuable medium for clinical investigation and nutritional status of human beings and animals [5]. In addition to the nutritional value, haemolymph of A. marginata cause agglutination of certain bacteria which are responsible for various ailments, including whooping cough. The high iron content is considered important in the treatment of anaemia [6]. It has also been reported to arrest constipation, restore vitality and stop bleeding when applied to a fresh cut. Previous biological studies reported that the acharan sulphate and mucin motifs isolated from A. marginata exhibited anti-tumor properties [7], and consistent blood glucose lowering effect [8] respectively. Recently the haemolymph of A. marginata has also been reported to have hepatoprotective effect and produce a dose dependent effect on haematological and biochemical parameters when administered to albino rats [3;9]. However, a literature survey has shown that there is no report on the bioactive constituents of the haemolymph thus the present investigation was carried out to identify active ingredients present in the Archachatina maginata haemolymph by (GC-MS) analysis

II. Materials And Methods

2.1 Snail Collection

African Giant Snails (*Achachatina maginata*) weighing 110-200g were bought from Kure market, Minna Niger state in September, 2014. They were housed in a ventilated container and fed with cucumbers and melon.

DOI: 10.9790/3008-10245964 www.iosrjournals.org 59 | Page

2.2 Haemolymph Collection

The haemolymph of *Archachatina maginata* was obtained as described by Bashir et al., [3]. The apex shell of the snails was opened; the haemolymph was drained into a clean conical flask and stored in the refrigerator.

2.3 Gas Chromatography-Mass Spectrometry (GC-MS)

GC-MS analysis was carried out on a GC-MS (Model: QP2010 PLUS Shimadzu, Japan) comprising a AOC-20i auto-sampler and gas chromatograph interfaced to a mass spectrometer (GC-MS) The instrument is equipped with a VF 5 ms fused silica capillary column of 30 m length, 0.25 mm diameter and 0.25 µm film thickness. The temperatures employed were; column oven temperature 80°C, Injection Temp 250°C at a pressure of 108.0 kPa, with total flow and column flow of 6.20 ml/min and 1.58 ml/min respectively. The linear velocity was 46.3 cm/sec and a purge flow of 3.0 ml/min. The GC program ion source and interface temperature were 200.00°C and 250.00°C respectively with solvent cut time of 2.50 min. The MS program starting time was 3.00min which ended at 30.00 min. with event time of 0.50 sec, scan speed of 1666 µl/sec, scan range 40-800u and an injection volume of 1 µl of the plant extract (split ratio 10:1). The total running time of GC-MS was 30 min. The relative percentage of the extract was expressed as percentage with peak area normalization.

2.4 Identification of the components

Interpretation on the mass spectrum was conducted using the database of National Institute Standard and Technology (NIST) having more than 62,000 patterns. The fragmentation pattern spectra of the unknown components were compared with those of known components stored in the NIST library. The relative percentage amount of each bio-component was calculated by comparing its average peak area to the total area. The name, molecular weight and structure of the components of the test materials were ascertained.

III. Results And Discussion

GC-MS chromatogram of the *Archachatina maginata* haemolymph (Fig. 1) showed seven peaks. The gas chromatogram shows the relative concentrations of various compounds getting eluted as a function of retention time. The heights of the peak indicate the relative concentrations of the components present in the haemolymph. The mass spectrometer analyzes the compounds eluted at different times to identify the nature and structure of the compounds. The large compound fragments into small compounds giving rise to appearance of peaks at different m/z ratios. These mass spectra are fingerprint of that compound which can be identified from the data library [10] On comparison of the mass spectra of the constituents with the NIST library twenty six (26) constituents including 7 ester, 7 fatty acid, 5 alcohol, 6 alkane and 1 phthalate were characterized and identified. The active principles with their retention time (RT), molecular formula, molecular weight (MW) and peak area (%) in the haemolymph are presented in Table 1.The mass spectrum and structure of the individual components are shown in Fig. 2. The first compound identified with less retention time (16.876s) was Methyl 14-methylpentadecanoate whereas 2-Methyloctadecane was the last compound which took longest retention time (25.254s) to identify. Literatures are scarce regarding the identified chemical constituents of *Archachatina maginata* haemolymph, however, The activities of some of the compound are given in table 2.The compound bioactivity prediction is based on Dr. Duke"s Phytochemical and Ethnobotanical Databases [11]

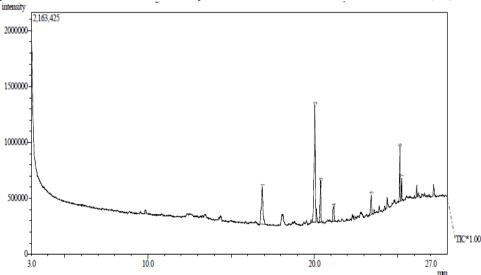


Figure 1.GC-MS Chromatogram of Archachatina maginata haemolymph

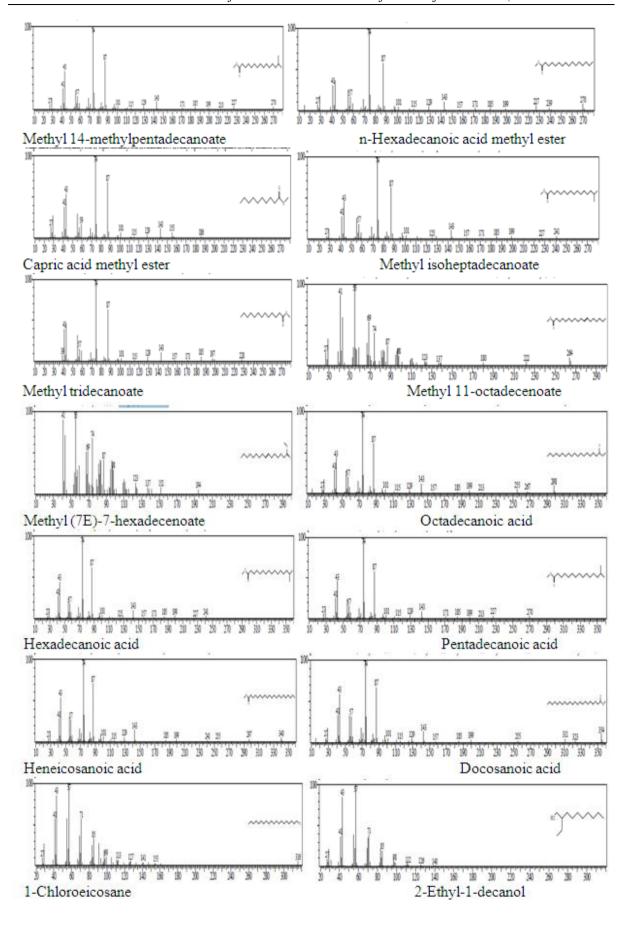
Table 1. Bio-active components identified in the Archachatina maginata haemolymph by GC-MS analysis

	Peak no	Retention (RT)(s)	Compound name	Nature of compound	Molecular formula (MF)	Molecular weight (MW-g/mol)	Peak area
1	1	16.876	Methyl 14- methylpentadecanoate	Ester	C17H34O2	270	19.21
2		16.876	n-Hexadecanoic acid methyl ester	Ester	C17H34O2	270	19.21
3		16.876	Capric acid methyl ester	Ester	C11H22O2	186	19.21
4		16.876	Methyl isoheptadecanoate	Ester	C18H36O2	284	19.21
5			Methyl tridecanoate	Ester	C14H28O2	228	19.21
6	2	20.041	Methyl 11-octadecenoate	Ester	C19H36O2	296	44.10
7		20.041	9-Octadecenoic acid	Fatty acid	C19H36O2	296	44.10
8		20.041	10-Octadecenoic acid	Fatty acid	C19H36O2	296	44.10
9		20.041	Methyl (7E)-7-hexadecenoate	Ester	C17H32O2	268	44.10
10	3	20.378	Octadecanoic acid	Fatty acid	C19H38O2	298	10.13
11		20.378	Hexadecanoic acid	Fatty acid	C18H36O2	284	10.13
12		20.378	Pentadecanoic acid	Fatty acid	C17H34O2	270	10.13
13		20.378	Heneicosanoic acid	Fatty acid	C22H44O2	340	10.13
14		20.378	Docosanoic acid/Behenic acid	Fatty acid	C23H46O2	354	10.13
15	4	21.149	1-Chloroeicosane	alkane	C20H41Cl	316	5.24
16		21.149	2-Ethyl-1-decanol	alcohol	C12H26O	186	5.24
17		21.149	1-Iodo-2-methylundecane	alkane	C12H25I	296	5.24
18		21.149	2-Butyl-1-octanol	alcohol	C12H26O	186	5.24
19	5	23.425	2-Methyloctadecane	alkane	C19H40	268	5.66
20		23.425	11,20-Di-n-decyltriacontane	alkane	C50H102	702	5.66
21		23.425	n-Nonadecane	alkane	C19H40	268	5.66
22	6	25.143	Bis(2-ethylhexyl) 3- nitrophthalate	Phthalate	C24H37NO6	435	11.21
23	7	25.254	2-Butyl-1-octanol	Alcohol	C12H26O	186	4.45
24		25.254	2-Hexyl-1-octanol	Alcohol	C14H30O	214	4.45
25		25.254	2-Hexyl-1-decanol	Alcohol	C16H34O	242	4.45
26		25.254	2-Methyloctadecane	Alkane	C19H40	268	4.45

Table 2. Activities of some identified compound in Archachatina maginata haemolymph

compound name	Activities			
n-Hexadecanoic acid methyl ester	Antioxidant, hypocholesterolemic, nematicide, hemolytic, 5-alpha reductase inhibitor			
Capric acid methyl ester	Calcium antagonist			
Methyl 11-octadecenoate	Allelopathic, pesticide			
9-Octadecenoic acid	Anti-inflammatory, Anti-alopecic, Anemiagenic, 5 reductase inhibitor, α-reductase inhibitor lubricant, Antitumour, Choleretic, Dermatitigenic, Immunostimulant, Anti-leucotriene-D4, Antiandrogenic, Lipoxygenase inhibitor, Allergenic, Flavour, Hypocholesterolemic, Insectifuge, Irritant, Percutaneo-stimulant, Perfumery and Propecic			
Hexadecanoic acid	Lubricant, antiandrogenic, antioxidant, 5- alpha-reductase inhibitor.12			
Pentadecanoic acid	Antioxidant			
Docosanoic acid/Behenic acid	Hair moisturizer			
1-Iodo-2-methylundecane	Sex hormones			
n-Nonadecane	Antimutagenic			

Sources: Dr. Duke"s Phytochemical and Ethnobotanical Databases [11].



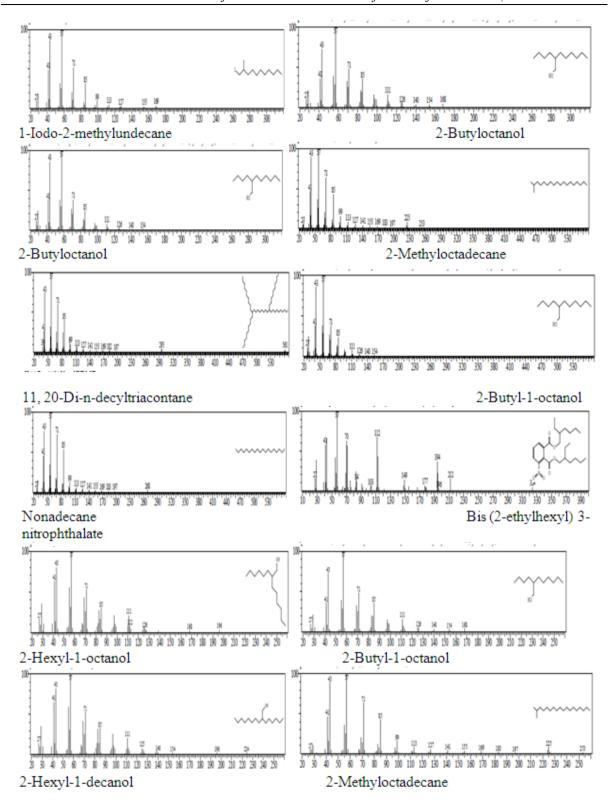


Figure 2; Mass fragmentation of the active components identified in the Archachatina maginata haemolymph

IV. Conclusion

In the present study twenty six chemical constituents have been identified from *Archachatina maginata* haemolymph by Gas Chromatogram- Mass spectrometry (GC-MS) analysis. The presence of various bioactive compounds justifies it uses for various ailments by traditional practitioners. However isolation of individual constituents and subjecting it to biological activity will definitely give fruitful results. It could be concluded that

DOI: 10.9790/3008-10245964 www.iosrjournals.org 63 | Page

Achachatina maginata haemolymph contains various bioactive compounds. So it is recommended as a wild animal of pharmaceutical importance.

Competing interest

The authors declare that they have no competing interest

ACKNOWLEDGEMENTS

Thanks are due to Adamu Mohammed of National Cereals Research Institute Zaria, Nigeria for help rendered during the GCMS analysis.

References

- [1]. Grace VMB, Manjamalai1 A, Yasaswini N, Aiswarya H, Antifungal, Antiinflammatory and GC MS Analysis Of Methanolic Extract Of Plectranthus Amboinicus LEAF.Int J Curr Pharm Res,2011Vol 3, Issue 2, 129136
- [2]. Sermakkani M, Thangapandian V.GC-MS Analysis of Cassia ItalicaLeaf Methanol Extract .Asian J Pharm Clin Res, Vol 5, Issue 2, 2012. 90-94
- [3]. Bashir L, Shittu OK, Prince CO, Asmau AN. and Aisha MI Evaluation of Antioxidant Activity of Giant African Snail (Achachatina maginata) Haemolymph in CCl₄- Induced Hepatotoxixity in albino Rats. Brit J Pharm Res.2015; 6(3): 141-154,
- [4]. Babalola OO, Akinsoyinu OA. Proximate, Mineral and Biochemical Evaluation of the Haemolymph of Growing Snails Fed Fresh Lettuce Waste, Whole Lettuce, Pawpaw Leaves and Cabbage Waste. A sian Journal of Agricultural Sciences 3(1): 1-4, 2011
- [5]. Cobbinah, J. R. (1992). Snail farming in West Africa a practical guide. Wageningen: Technical Centre for Agricultural and Rural Co-operation. British Crop Protection Council Monograph 11: 49-55.
- [6]. Lee, W. G., Mark, A. F. and Wilson, J. B. (2003). Ecotypic differentiation in the ultramafic flora of the South Island, New Zealand. New Zealand Journal of Botany 21: 141-156.
- [7]. Cooling, V. (2005). Risk Assessment of the Giant African Snail Bowdich in New Zealand. LPSC 7700 Integrative Report
- [8]. Adikwu MU. Evaluation of snail mucin mitifs as rectal absorption enhancer for insulin in non-diabetic rat models. Bio Pharm Bull. 2005;28(9): 1801-4
- [9]. Bashir L, Shittu OK, Busari MB, Sani S, Aisha MI-Safety Evaluation of Giant African land Snails (Achachatina marginata)
 Haemolymph on Hematological and Biochemical Parameters of Albino Rats. Journal of advances in medical and pharmaceutical sciences (in press)
- [10]. Janakiraman N, Johnson M, Sahaya Sathish S/.(2012). GC-MS analysis of bioactive constituents of Peristrophe bicalyculata (Retz.) Nees. (Acanthaceae). Asian Pacific Journal of Tropical Biomedicine (2012)S46-S49
- [11]. Duke"s Phytochemical and Ethnobotanical Database; 2014. Available: http://www.ars-grin.gov/cgibin/ duke/ethnobot/ Assessed 16th Jun. 2014

DOI: 10.9790/3008-10245964 www.iosrjournals.org 64 | Page