

## Extraction and Characterization Some Fatty Acid of the Seeds Oil from *Albizialebbeck (L.) Benth*

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### Abstract

### Original Research Article

*Albizialebbeck(L.) Benth.* A member of Fabaceae family, have numerous medicinal and pharmacological properties, the plant was collected from Elfitahab areas in Omdurman city. The fruit pulp of *Albizialebbeck(L.) Benth.* was subjected to continuous and successive soxhlet extraction with n-Hexane . The oil extract was screened with antimicrobial activity and analysis by GC/MS technique to appeared 19 components.

**Keywords:** *Albizialebbeck(L.) Benth*, GC/MS, Antimicrobial activity, Soxhlet.

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## INTRODUCTION

*Albizialebbeck(L.) Benth* tree it is belonging to the family Fabaceae member of the subfamily Mimosae It is widely available plant in the tropical and subtropical Asia and Africa [1]. It is cultivated in many parts of Sudan as shade tree in front of houses, farmlands, along roadsides, along rivers, and as an ornamental plant in gardens due to its pleasant appearance. It evergreen tree with height about 18-30m with a trunk 50cm to 1m in diameter and the bark is gray colour [2] with glabrous young shoots [3]. Leaves are evenly bi pinnate and the leaflets are in 5-9 pairs, 2.5-5.0 cm long, broadly oblong and pale green [1] flowers are to greenish-yellow ,fragrant in globose umbellate heads 2-3.8cm diameter, stamens are longer than the corolla [2]. Pods are 10-30 cm long, 2-4.5 cm wide, thin, linear oblong, smooth, green turning yellow-brown on maturity, They have 4-12 yellowish brown seeds [4, 5]. The plant have numerous medicinal and pharmacological properties, different parts of this plant have lots of useful properties, the roots are used in ophthalmia, alleviate spasms, rat bite, stimulate the cardiovascular system, hemicranias, anticancer and spermicidal properties, antifungal activity [2, 3] the bark is used for treatments of skin disease, relives toothache, hemorrhoid, leprosy, piles, deafness, cooling, alexiteric, pimples, itching, anthelmintic, syphilis, cures diseases of blood, leucoderma, itching, paralysis, cures diseases of blood, leucoderma, itching and body weakness, bubbles, boils, scabies, Inflammation,

antidiabetic, bronchitis, over the top sweat, excessive perspiration, heaps, After drying and pounding, it is used as a soap substitute [3, 6, 7, 2]. The leaves of this tree are good for syphilis, ophthalmic diseases, night blindness, nootropic activity, syphilis and ulcer, anxiolytic activity, cold, cough, respiratory disorder [3, 2]. The leaves are also used as cattle fodder, mulch, and excrement because of high nitrogen contents [3, 6] Leaves are also used as soap substitute after being dried [7]. The flowers are used in treatment of ophthalmia, aphrodisiac, emollient, night blindness, maturant, psychological disorders, spermatorrhea. insomnia and warts, Flowers also are applied as cataplasm on furuncles and used for retention of seminal fluid [3, 8].

The pods extract is reckoned have antidiabetic, antiprotozoal, antidiabetic, and anticancer properties and also show antispermatogenic effect [2, 3] and possesses a wide antibacterial property against *E. coli* and antiamebic activity against *Entamoeba histolytic* [9]. The Seeds have been used for a variety of therapeutic purposes such as cure piles, can treat diabetes, anti-tumor, diarrhea, scrofulous swelling, aphrodisiac, brain tonic, gonorrhoea and tuberculous glands [2, 10]. And have constipating properties, astringent. Are also used for treating hemorrhoid, it is believed that the extract of seed covers have anti protozoal properties [7], The seeds oil is applied topically to cure leucoderma [3]. The oliage is commonly used as fodder in India for cattle, and in Sudan, goats eat fallen leaves and flowers [11].

## MATERIALS AND METHODS

### Plant Material

*Albizialebbeck*(L.) Benth seeds were collected from faculty of agriculture Omdurman Islamic University Elfetehab area Omdurman city, Khartoum State (Sudan), during April 2018. The plant is authenticated by the Department of Biology and Chemistry faculty of Education, Omdurman Islamic University. The seeds of *Albizialebbeck*(L.) Benth were

manually separated from the fruits, dried at room temperature at 8 days, 500 g were ground by an electric mixer until a fine powder is obtained 367 g and stored at room temperature until further use.

### Test Organisms

*Albizialebbeck*(L.) Benth oil was screened for antibacterial and antifungal activities using the standard microorganisms shown in Table-1.

**Table-1: Test organisms**

Ser. No	Micro organism	Abb.	Type
1	<i>Bacillus subtilis</i>	<i>B.s</i>	G+ve
2	<i>Staphylococcus aureus</i>	<i>S.a</i>	G+ve
3	<i>Pseudomonas aeruginosa</i>	<i>P.a</i>	G-ve
4	<i>Escherichia coli</i>	<i>E.c</i>	G-ve
5	<i>Candida albicans</i>	<i>C.a</i>	Fungus

### Extraction of oil

Powdered shade-dried seeds of *Albizialebbeck*(L.) Benth s(500g) were exhaustively extracted with n-hexane at room temperature. The solvent was removed *in vacuo* and the oil was kept in the fridge at 4oC for further manipulation.

### Esterification of oil

A Methanolic solution of sodium hydroxide was prepared by dissolving (2g) of sodium hydroxide in 100ml methanol. A stock solution of methanolic sulphuric acid was prepared by mixing (1ml) of concentrated sulphuric acid with (99ml) methanol.

The oil (2ml) was placed in a test tube and 7ml of alcoholic sodium hydroxide were added followed by 7ml of alcoholic sulphuric acid. The tube was stoppered and shaken vigorously for five minutes and then left overnight.(2ml) of supersaturated sodium chloride were added, then (2ml) of normal hexane were added and the tube was vigorously shaken for five minutes .The hexane layer was then separated. (5µl) of the hexane extract were mixed with (5ml) diethyl ether. The solution was filtered and the filtrate(1µl) was injected in the GC-MS vial.

### Instruments

Gas Chromatography coupled with Mass Spectrometry Analysis was conducted on a Shimadzo GC-MS-QP2010 Ultra instrument equipped with a split-splitless injector and with a RTX-5MS column (30m, length; 0.25mm diameter; 0.25 µm, thickness). The GC oven was set to a temperature range of 150 to 3000C with 60C/min, and a solvent delay of 7 min. The injector temperature was maintained at 300 °C. The carrier gas of the sample was helium at 1.0 mL/min and the sample was injected in the splitless mode. The MS conditions were the followings: ionization energy, 70 eV; electronic impact ion source temperature, 200<sup>0</sup>C; interface temperature, 25<sup>0</sup>C; scan rate 1.6 scan/s; mass, 40-600 amu. For the identification the

compounds the mass spectra of the samples were compared with determined by the chromatographic method with the help of NIST and WILEY library.

### Antimicrobial Assay

One gram of the oil was weighed and dissolved in 10ml of DMSO to obtain a concentration of 100mg/ml. This was the initial concentration of the oil used to check the antimicrobial activities. Diffusion method was the method used for screening the oil. Mueller Hinton and Sabouraud dextrose agars were the media used as the growth media for the bacteria and the fungus respectively. The media were prepared according to the manufacturer's instructions, sterilized at 121oC for 15 minutes, poured into sterile Petri dishes and were allowed to cool and solidify. The sterilized media were sealed with 0.1ml of the standard inoculums of the test microbe (Mueller Hinton agar was sealed with the bacteria and Sabouraud dextrose agar sealed with the fungus). The inoculums were spread over the surface of the medium by the use of a sterile swab. By the use of a standard cork borer of 6mm in diameters, a well was cut at the centre of each inoculated medium. Serial dilutions of the oil (0.1ml) were then introduced into the well on the inoculated medium. Incubation of the inoculated medium was made at 37oC for 24 hours for the bacteria and at 30oC and for 4 days for the fungus. After incubation each plate of the medium was observed for the growth inhibition zone. The zone was measured with a transparent ruler and the results were recorded in millimeters.

## RESULTS AND DISCUSSION

### Constituents of oil

The GC-MS spectrum of the n-hexane extract of oil is shown in the Figure 1. Total of 19 compounds present in the n-hexane extract were determined by the chromatographic method with the help of NIST and WILEY library as shown in Table-2.

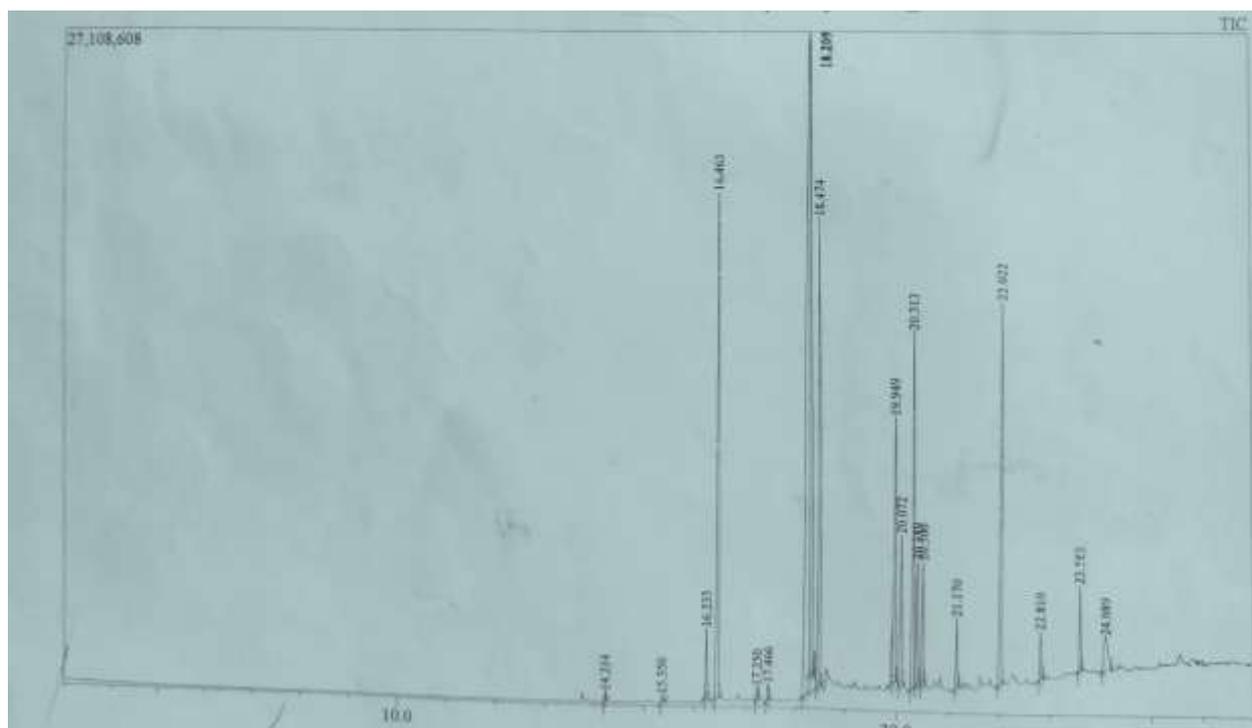


Fig-1: *Albizialebeck(L.)* Benth seeds oil Chromatogram

Table-2: Fatty acids constituent of *Albizialebeck(L.)* Benth seeds oil

Peak No.	Name	R.Time	Area %
1	Methyl tetradecanoate	14.214	0.17
2	Pentadecanoic acid ,methyl ester	15.350	0.12
3	9-Hexadecenoic acid, methyl ester,(Z)	16.235	1.11
4	Hexadecenoic acid, methyl ester, (Z).	16.463	12.32
5	Cis-10- heptadecenoic acid, methyl ester	17.250	0.22
6	heptadecenoic acid, methyl ester	17.466	0.27
7	9,12-Octadecadienoic acid (Z,Z),methyl ester	18.250	29.09
8	9-Octadecenoic acid (Z),methyl ester	18.295	13.03
9	Methyl Stearate	18.474	9.72
10	E,E,Z-1,3,12-Nonadecatriene-5,14-diol	19.949	5.39
11	Oxaraneoctanoic acid,3-octyl,methyl ester	20.017	3.58
12	Eicosenoic acid, methyl ester	20.313	7.31
13	PGTI, methyl ester	20.389	2.21
14	Mehtyl-13,16-Docosandienoate	20.500	2.24
15	Heneicosanoic acid, methyl ester	21.170	1.19
16	Docosenoic acid, methyl ester	22.022	8.19
17	Tricosanoic acid, methyl ester	22.810	0.88
18	Tetracosanoic acid, methyl ester	23.583	1.44
19	Gamma.sitosterol	24.89	1.53
Total of components			100%
Total Saturated fatty acids			17.10
Total Unsaturated fatty acids 82.90			
- Total of Mono unsaturated fatty acids 46.18			
- Total of Poly unsaturated fatty acids 36.72			

GC/MS analysis of *Albizialebeck(L.)* Benth seeds oil provided a rich source of unsaturated fatty acids(82.90%), important among which is linoleic acid with hypocholesterolemic effect, implying a cardioprotective property, then the amount of saturated fatty acids is (17.10%), but unsaturated fatty acids contains monounsaturated fatty acids (46.18 %) and

polyunsaturated fatty acids (36.72%). The fatty acids acquire high percentage in *Albizialebeck(L.)* Benth seeds oil were 9,12-Octadecadienoic acid (29.09%), 9-Octadecenoic acid (13.03%), Hexadecenoic acid, methyl ester, (Z).(12.31), Methyl Stearate (9.72), Docosenoic acid, methyl ester (8.19%), Eicosenoic acid, methyl ester (7.31), E,E,Z-1,3,12-Nonadecatriene-

5,14-diol d (5.39%), Oxaraneoctanoic acid, 3-octyl, methyl ester (3.58), other fatty acids which were a counted in percentage of the total fatty acids is modest amount.

### Antimicrobial Activity

The oil was screened for antimicrobial activity against five standard microorganisms. The average of the diameters of the growth inhibition zones are shown in Table-3. The results were interpreted in terms of the

commonly used terms (>9mm: inactive; 9-12mm: partially active; 13-18mm: active;<18mm:very active). Table-3 represented the antimicrobial activity of standard antibacterial and antifungal chemotherapeutic agents against standard bacteria and fungi respectively. The oil showed activity against *Pseudomonas aeruginosa* and partial activity against *Escherichia coli*, but it is inactive against *Staphylococcus aureus*, *Bacillus subtilis*, *Staphylococcus aureus* and the fungi *Candida albicans*.

**Table-3: Antimicrobial activity of standard drugs and *Albizialebbeck(L.) Benth seeds oil*.**

Name of compound	Conc.	Minimal inhibition concentration ( $\mu\text{g mL}^{-1}$ )				
		Gram-negative		Gram-positive		Fungal species
		P.a	E.c.	B.s.	S.a	C.a
<i>Albizialebbeck(L.) Benth seeds oil</i>	100	15	11	-	-	-
Gentamycin	40	21	22	25	19	-
	20	15	18	22	18	-
	10	12	15	17	14	-
Nystatin	-	-	-	-	-	100

### CONCLUSIONS

In the present study, the authors reported the chemical constituents and biological activities of the *Albizialebbeck(L.) Benth seeds oil*. The different part of plant are showed Biological activities that is this plant display a multidisciplinary usage in treating several diseases. Further separation and identification of compound present in it may give new biologically active compounds, which can be used as lead compounds in future.

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