

Structure Determination of Compounds from The Stem of *Equisetum Ramosissimum*

Abstract

This study was carried out to isolate the major constituents of stem of *Equisetum ramosissimum*. The DME extract was isolated by column chromatography to give one pure compound. The compound PE-2 (11 mg) was identified as Isophthalic acid 1-(5-methyl-hexyl) ester 3-pentyl ester. The structures of the compounds were elucidated by means of ¹H-NMR, ¹³C-NMR and DEPT-135 spectral data and comparison with literature reports.

Keywords: Chemical constituents; some major compounds from stem of *Equisetum ramosissimum*

Introduction

Plants are important in human's life and fulfill his every day needs. Different plant parts like root, stem, flower, leaves, fruit, twigs and modified plant organs are used as food, Shelter, clothing, transportation, fertilizer, flavors, fragrances, ornamental, medicine, cosmetic, etc. throughout the ages of humans. According to some observation some animals like chimpanzees utilize a number of plant species for medicinal use [1].

The discovery of plants that serve as medicine (medicinal plant) in different parts of the world is important to agriculture and medicine sectors, in establishment of new directions towards propagation of alternative medicinal plant that offer better economic and social benefits. The ancient civilizations such as Chinese, Egyptian, Indians, Greek, and North Africans provide written evidence for the use of natural products for treatment of various diseases [2].

The use of medicinal plants, still play a vital role to cover the basic health needs in the developing countries like Africa, in many African countries traditional medicines from medicinal plants are sold in marketplace or prescribed by traditional healer. For example, mandrake was prescribed for pain relief, turmeric possesses blood clotting properties, roots of the endive plant were used for treatment of gall bladder disorders, and raw garlic was prescribed for circulatory disorders [2].

The world health organization (WHO) reported that 80% of the world's populations depend on traditional medicine and a major part of traditional therapies involve the use of medicinal plant extracts or their active constituents [3].

Medicinal plants used traditionally throughout the world to treat many illnesses like malaria, diabetes, respiratory and urinary tract infections, cough, fever, diarrhea, abdominal pains, pneumonia, conjunctivitis, oral and tooth wounds etc. and also used for birthcontrol and psychic problems and often exhibit a wide range of biological and pharmacological activities, such as anti-bacterial and anti-fungal properties. Due to the need for development of new compounds with better pharmacological activities, dependence on medicinal plants grew increasingly as scientists continuously exploited them for isolation of bioactive compounds [4]

On this regard *Equisetum ramosissimum* is a medicinal plant which is popularly called “horsetail”, and locally called “riga bofa” is used by traditional medicine practitioners as wound treatment, toothache, treating urinary tract infection, cardiovascular diseases, respiratory tract infection and medical skin conditions [5].

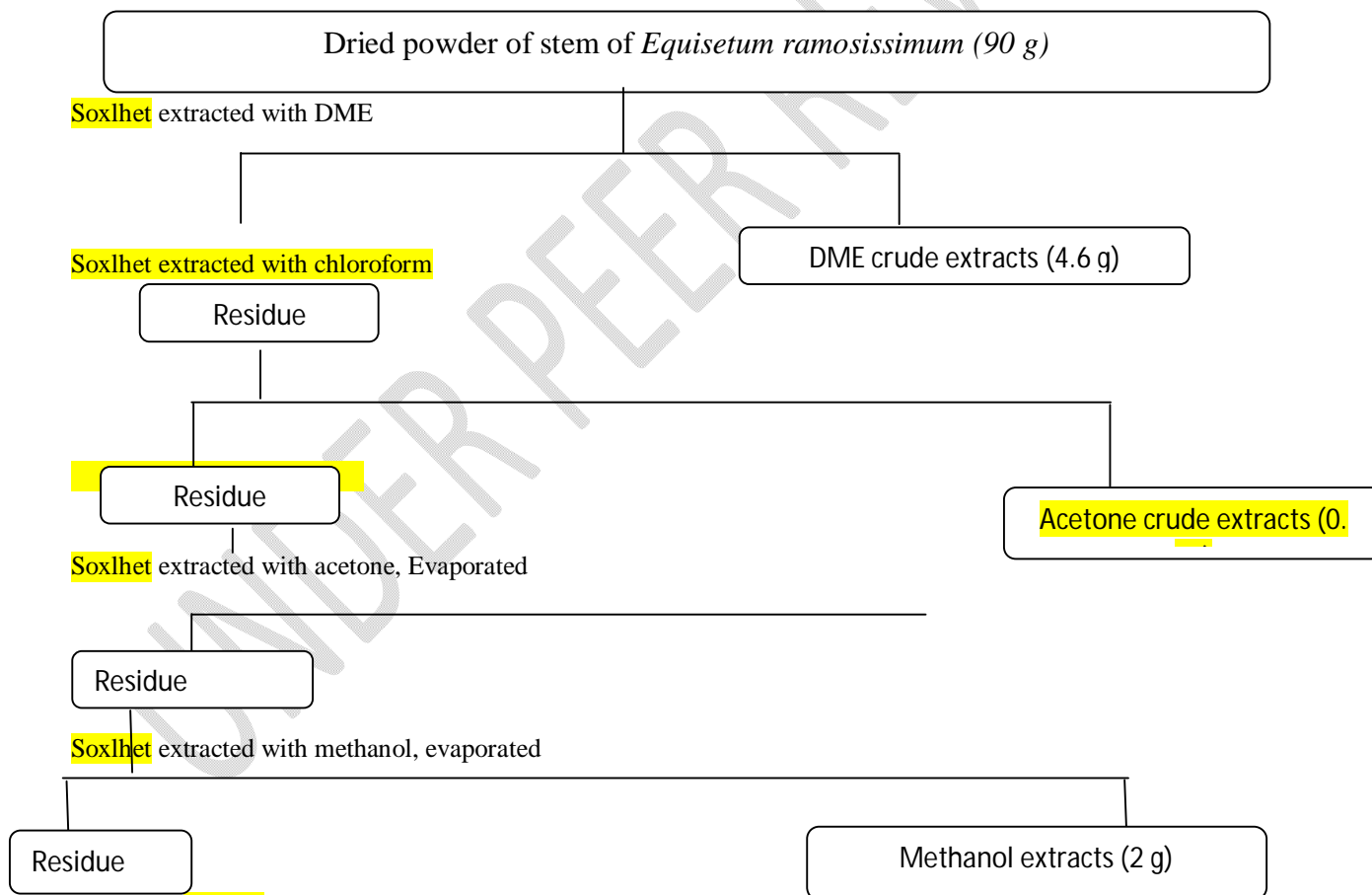
Materials and Method

Plant Material Collection and Identification

The Stems of *Equisetum ramosissimum*. Local name (*Riga boofaa*) in Afaan Oromo was collected from Amuru village, Horro Guduru, Welega Zone, Oromia Region, which is 383 km west of Addis Ababa. The plant was identified by prof. Legesse Negash and specimen was deposited at the National Herbarium (Voucher Diriba Borena 002/2015) the Department of Biology, Addis Ababa University.

Experimental procedures Extraction

90 g (divided into three parts) of powdered stem of *Equisetum ramosissimum* were extracted by using dimethyl ether, chloroform, acetone and methanol (350 ml for each) in Soxhlet apparatus [16]. The extraction procedure is given in the next scheme.



Scheme 1 Soxhlet extraction procedure

Isolation of Compounds

According to the TLC analysis of petroleum ether: ethyl acetate for good separation of ether extract was 6:4 and methanol: ethyl acetate was 8:2. Depending on this ratio of solvent selected column chromatography was

packed with petroleum ether after absorbing the dimethyl ether extract with silica gel and concentrated on rotary evaporator. The extract was applied on column and eluted with increasing polarity of petroleum ether/EtOAC solvent mixtures. Elution of the column by PE: EtOAC is given in the next figure.



Figure 10 Isolation of DME extract into fractions

Spectroscopic analysis

Pure fractions from column chromatography were characterized by using IR, UV and nuclear magnetic resonance ($^1\text{H-NMR}$, $^{13}\text{C-NMR}$, DEPT) and the spectra were recorded in CDCl_3 and DMSO-d_6 with **Tetramethylsilane** (TMS) as internal standard. Complete structure determination was achieved by comparing the IR and NMR data obtained with that in literature.

Isolation of compounds from PE-EtOAC Extract of stem of *Equisetum ramosissimum*

Compound **PE-2** (11 mg) was isolated using CC from non-polar fraction as blue black solid. TLC analysis by PE: EtOAC (6:4) showed a single spot with $R_f = 0.7$ staining red under UV light. This compound is most likely one of a non-polar and $^1\text{H NMR}$, $^{13}\text{C NMR}$ and DEPT-135 spectral data of the compound is given in table 1 below.

Table 1 $^1\text{H NMR}$, $^{13}\text{C NMR}$ and DEPT-135 for PE-2

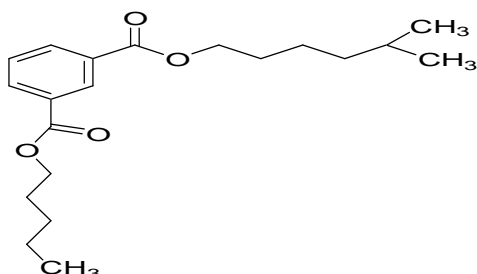
No	^1H δ (ppm)	^{13}C δ (ppm)	DEPT-135 δ (ppm)	Remark
1		167.00		COO-
2		132.37		
3	7.763	130.94	130.95	CH=
4	7.748	128.82	128.82	CH=
5	4.189	68.19	68.20	CH ₂ -O
6	2.046	31.94	31.94	CH ₂
7	1.019	30.36	30.36	CH ₂
8	1.608	29.71	29.71	CH ₂
9	1.831	28.83	28.93	CH
10		27.73	27.73	
11	1.277	23.75	23.75	CH ₂
12	1.002	22.71	22.71	CH ₃ -CH
13		19.39	19.39	
14	0.902	14.14	14.16	CH ₃

The $^1\text{H NMR}$ spectrum (Appendix A and, Table 1) of the compound showed peaks δ 0.902 (6H, and δ integrating for two methyl protons. A methylene signals appeared at δ 1.277 and δ 1.608 each integrating for two protons each. Other methylene signals appeared at δ 1.019 and 2.046 each integrating for two protons each.

Ox methylene signals appeared at δ 4.189. Aliphatic methine proton peaks appeared at δ 1.831. Furthermore, the two overlapped signal appeared at assigned for δ 7.748 the two olefinic methane protons. One signal is observed δ 7.763 for olefinic methine proton.

The ^{13}C NMR and DEPT-135 (Appendix B, C and Table 1) indicated that compound 3 has 20 carbon atoms. The spectra showed two aliphatic methyl carbons at δ 22.71 and one aliphatic methyl carbon at δ 14.14. Two aliphatic methylene carbons at δ 23.75 and 29.71, two methylene carbons at 31.94, two methylene carbon at δ 30.36., two other methylene carbon at δ 66.20. Two ox methylene carbon atoms appeared at δ 68.19 one aliphatic methine at δ 28.83, two olefinic methine at δ 128.83., one methine at δ 130.94, and one two olefinic quaternary carbon at δ 132.37, two ester carbonyl carbons at δ 167.00.

Based on the above NMR data the next structure was proposed for the compound PE-2.



Isophthalic acid 1-(5-methyl-hexyl) ester 3-pentyl ester

Figure 2 Tentative structure of compound PE-2

Conclusion

The dried powder of stem of *Equisetum ramosissimum* was extracted with DME, chloroform, acetone and methanol successively. The DME extract was isolated by column chromatography to give one pure compound. The chemical structure was characterized on the basis of NMR spectral data, including ^1H -NMR and ^{13}C -NMR in comparison with literature values. The compound PE-2 (11 mg) was identified as Isophthalic acid 1-(5-methyl-hexyl) ester 3-pentyl ester.

Acknowledgement

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NOTE:

The study highlights the efficacy of " traditional medicine " which is an ancient tradition, used in some parts of India. This ancient concept should be carefully evaluated in the light of modern medical science and can be utilized partially if found suitable.

References

1. Huffman M. and Wrangham R. Diversity of medicinal plants use by wild Chimpanzee behavioral diversity. *Harvard University press*.**1993**, **27**, 1-14.
2. Phillipson J. Phytochemistry and Medicinal Plants. *Phytochemistry*. **2001**, **56**, 237–243.
3. Herbal G. summary of WHO guidelines for assessment of herbal medicine. *World health organization*. **1993**, 28, 28-43
4. Saliu B., Usman L., Sani A., Muhammad N. and Akolade J. Chemical composition and antibacterial (oral isolates) activity of leaf essential oil of *Ocimum gratissimum* L. grown in north central Nigeria. *International Journal of Current Research*. **2011**, **33**, 022-028.
5. Ejele A., Iwu I., Enenebeaku C., Ukiwe L. and Okolue B. Bioassay-Guided Isolation, Purification and Partial Characterization of Antimicrobial Compound from Basic Metabolite of *Garcinia Kola*. *Journal of emerging trends in engineering and applied sciences* .**2012**, 3, 668-672.
6. Ivana D., Milan S., Olgica D., Marina D., Ljiljana R. and Aleksandar M. Great Horsetail (*Equisetum telmateia* Ehrh), Active substances content and biological effects. *ExcliJournal* .**2012**, 11,59-67.