

Original Research Article

Phytochemical Screening of Root, Stem and Leave

Extracts of *Terminalia avicennoides*

ABSTRACT

The phytochemical screening of *Terminalia avicennoides* was carried out using qualitative method to determine the bioactive compounds present in the plant root, stem and leave extracts. Cooled Maceration method was used for the extraction. Hundred grams (100g) of each powder was soaked in 1000ml of distilled water, allowed to stand for 5hours. The suspension was agitated after 30 minutes. The filtrate was thereafter separated from residue using No. 1 Whatman filter paper and concentrated using rotary evaporator. The crude extracts were separately kept in a screw capped bottle for further research. The bioactive compound in the plants were detected using AOAC method. The result revealed that alkaloid, flavonoid, tannin, saponins, phenol and glycoside were detected in the plants while steroid was not detected in the plants. Therefore, the presence of these phytocompounds is an indicative that the plant is medicinal and it can be used for the treatment of bacterial and fungal infections.

Key words: Phytoscreening, *Terminalia avicennoides*

1.0 INTRODUCTION

Terminilia avicennoides also known as Combretaceae grow in the savannah region of West Africa (Mann, *et al.*, 2007). It is called “baushe” in Hausa and “Ungbo” in Adara. The plant is used for the treatment of burn wound infection in humans by the Adara people of North central of Nigeria. The use of the root necesacitated this works to determine the phytochemical composition of the entire plant parts.

The use of medicinal plants in the treatment of diseases has gained popularity and generated special interest in recent times. Herbal preparations are increasingly being used in both human and animal healthcare systems. Diarrhea is one of the common clinical signs of gastrointestinal disorders caused by both infectious and non-infectious agents leading to serious human and livestock debilitating condition. Herbal medicine has long been recognized as one of the oldest sources of remedies used by humans according (Mann *et al.*,

30 2009). A lot of people in both undeveloped and developing countries still rely on medicinal
31 plants for their daily healthcare needs, in spite of the advancement in modern medicine
32 (Passalacqua, *et al.*, 2006). Different traditional healing practices globally are designed for
33 either therapeutic or prophylactic use in human or animal diseases. A number of studies
34 carried out in Africa, Asia, Europe, Latin America and North America show that plants are
35 routinely used as remedy for animal diseases. Historically, it is documented that humans
36 make use of the same herbal preparations that they use to treat their sick animals. In Nigeria,
37 farmers are known to treat animal diseases with herbs before the advent of modern
38 medicine. Traditional medical and veterinary practices remain relevant and vital in many
39 areas in Nigeria (Ojewole, 2004).

40 Modern drugs are derivatives of medicinal plants (Abdullahi, 2012). It is now believed that
41 nature has given the cure of every disease in one way or the other. Plants have been known to
42 relieve various diseases worldwide. Plant-derived substances have recently become of great
43 interest owing to their versatile applications. Today, plant materials play a crucial role in the
44 health sector and many studies revealed plants are good sources of antimicrobials (Mann,
45 2012). Medicinal plants are the richest bio-resource of drugs, traditional systems of medicine,
46 modern medicines, nutraceuticals, food supplements, folk medicines, pharmaceutical
47 intermediates and chemical entities for synthetic drugs (Ncube *et al.*, 2008). The knowledge
48 of the phytochemicals present in extracts is a merit for area of specialization. The correct
49 identification of the herbal material and the active constituents is crucial to quality control,
50 safety, efficacy, acceptability and possible integration into the national healthcare system of
51 herbal remedies. This study was thus carried out to identify bioactive compounds present in
52 *Terminalia avicennoides* extracts, which also may be responsible for its antibacterial activity.

53 **2.0 MATERIALS AND METHOD**

54 **2.1 Collection, identification and authentication of the plant**

55 The fresh root, stem and leaves of *Terminalia avicennoides* was collected from the open field
56 at Doka village in Doka District along Kaduna-Abuja express way, Kachia Local
57 Government Area, Kaduna State, Nigeria. It was identified at the Department of Botany,
58 Faculty of Science, Ahmadu Bello University Zaria, with the herbarium number 900239.

59 **2.2 Preparation of the plant material**

60 The root, stem and leaf *Terminilia avicenoides* were collected in July, 2017. The grey bark
 61 was cleared. The inner bark was air-dried and later pulverized with the aid of mortar and
 62 pestle into powder form. The powder was packed into a clean plastic container with screw
 63 cap for subsequent bench work.

64 2.3 Extraction

65 Hundred gram of each powder (root, stem and leaf) was soaked in litre of distilled water and
 66 allowed to stand for five hours and was agitated after 30 minutes, after which the suspension
 67 was macerated. The extracts were successively extracted using aqueous solvent. With the
 68 help of vacuum evaporator, all the plant extracts were concentrated. Each extract was dried
 69 using hot-air oven at low temperature to a constant mass for further research.

70 2.4 Phytochemical analysis

71 The method of Trease and Elvans, (1996) was applied for the determination of the presence
 72 of phytochemicals. The filtrate obtained from each extraction was tested for alkaloids,
 73 saponins, flavonoids, phenol, steroids, tannins and glycosides.

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75 3.0 RESULT

76 The phytochemical analysis of aqueous extract of *T. avicenoides* revealed the presence of
 77 alkaloid, flavonoid, tannin, saponins and glycoside were detected while steroid was not
 78 detected in the plants.

79 **TABLE 1: Phytochemical Constituents of Root, Stem and Leaf Extracts of *T.***
 80 ***avicenoides***

81	PHYTOCOMPOUNDS	ROOT	STEM	LEAF
82	Alkaloids	+	+	+
83	Flavonoids	+	+	+
84	Tannins	+	+	+
85	Saponins	+	+	+
86	Steroids	-	-	-
87	Phenol	+	+	+
88	Glycosides	+	+	+

89 **Key: + Detected, - Not Detected.**

90 **4.0 DISCUSSION**

91 From Table 1, the phytochemical constituents of the plant extracts ‘root, stem and leaf’
92 showed the presence of alkaloids, flavonoids, tannins, saponins, glycosides and phenol.
93 Brantner and Grein, (2004) also carried out the same research and detected the presence of
94 alkaloids, tannins, saponins and using ethanol. This result indicates that the parts of the plants
95 have active ingredients responsible for antimicrobial activity. The presence of these
96 secondary compounds makes the plants fits or good for the treatment of bacterial and fungal
97 infections because most therapeutic effects of medicinal plants are traced to the plant
98 constituents and the medicinal actions of these constituents are unique to particular species or
99 family (Cowan, 1999). Also, from the result it can be deduced that only polar and
100 moderately polar phytocompounds are presence which may be due to the fact that, the
101 extraction was done using polar solvent (distilled water). It is possible that this plant may
102 have high antimicrobial activity due to the presence of these metabolites. Further study can
103 be done to separate the individual metabolites to test their antimicrobial activity against some
104 pathogenic bacteria to determine their potency.

105 **5.0 CONCLUSION AND RECOMMENDATION**

106 Phytochemical composition of the root, stem and leaf extracts of the *T. avicennoideis* indicate
107 the presence of six active constituents. The presence of these phytocompounds is an
108 indicative that the plant is medicine and it can be used for the treatment of bacterial and
109 fungal infections. Other solvents such as ethanol, methanol, and petroleum ether can be used
110 for extraction so that other phytocompounds can be detected. Further investigation,
111 purification and determination of these promising constituents can be done to assay their
112 antimicrobial activity.

113 **REFERENCE**

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