



Antioxidant and acute toxicity of stem extracts of the *Ficus iteophylla*

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Abstract

The aims of this Study is to evaluate the Antioxidant activity and acute toxicity of the extracts of *Ficus iteophylla* by reactions with 1, 1-diphenyl-2-picrylhydrazyl radical (DPPH) and method developed by Lork 1983 respectively. Stem of *Ficus iteophylla* was collected, air dried, pulverized to fine powdered and sequentially extracted using acetone, methanol and water in order of increasing polarity. The result show strong radical scavenging activity against DPPH for all the extracts when compared with Vit. C. The LD50 of 316mg/kg was calculated for all the three extras and the values were found to be within the practically toxic range and therefore care should be taken when using the plants in traditional medicine.

Keywords: antioxidant, acute toxicity and *Ficus iteophylla*

Introduction

Medicinal plants have long being used by man for treatment of various illnesses. According to world health organization (2001), 80% of the world population uses traditional medicine for the treatment of diseases. The number of people using traditional medicine is much higher in developing countries. World health organization estimated that 90% of the people leaving in developing countries depend on trational medicine for their primary health care need (WHO, 2002) [16]. Medicinal plants are widely accepted because they are safer, cheaper readily available than synthetic drugs (Ayonde and Adebire 2008). Despite the general assumption that medicinal plants are safer than synthetic drugs, there are many stuties that reported mutagenic, Cercinogenic and Toxic properties of some medicinal plants (Deciga *et al.*, 2007, Ferreira ICFS *et al.*, 1999 and Mohd-Fuat AR *et al.*, 2007) [3, 5, 13]. The term antioxidant refers to the activity of numerous vitamins, minerals, and other phytochemical to protect against the damage cause by reactive oxygen species (Dula 2018) [4]. The reactive oxygen species are involved in various physiological process diseases such as ageing, cancer, diabetes and atherosclerosis (Priyanka p, 2011) [15]. The antioxidant activities of medicinal plants are due to the presence of flavones, isoflavone, flavooids, anthocynin, catechin, isocatechin, phenolic compounds and tannins (Killedar 2013) [7]. Plants or drugs must be ensure to be safe before the can be uses as medicine. A key stage in ensuring the safety of drugs is to conduct a toxicity test in an appropriate animal. Acute toxicity studies is one of the procedure for toxicity test that are in use. The main aim of this study was to evaluate the antioxidant of the stem extract of *Ficus iteophylla* and also to evaluate it acute toxicity before it can be recommended for application that are important for public.

Material and Method

Collection of Plant Material

Ficus iteophylla are available at Hadejia Nguru wet land and it was collected from Gaffa area of the wet land. The plant was identified by the taxonomist at the herbarium of Ahmadu Bello University Zaria and is deposited with voucher number 2504. It was wash thoroughly, air dried and grounded to fine powdered.

Extraction

100g of the fine grounded plants was successively extracted using Acetone (1000ml), methanol (1000ml) and water (1000ml) for 48 hours and evaporated to formed acetone extract, methanol extract and water extract.

Experimental Animal

Experiment was performed using healthy swiss albino mice of both sexes. They were purchase from the Department of Pharmacology Aminu Kano Teaching Hospital, Bayero University. Kano.

Antioxidant Assay

The antioxidant activity of the extracts and standards were measured on the basis of the radical scavenging effect of the stable 1, 1-diphenylpicrylhydrazyl (DPPH) free radical activity method modified by Braca *et al.*, 2002. The working solutions of the of the extracts were prepared in methanol, Ascorbic acids was used as a standard in 0.01g/ml. 0.004% of DPPH was prepared in methanol and 2 ml of the sample was mixed with 2 ml of this solution and standard solution separately. These solution mixtures were kept in dark for 30 minutes and optical activity was measured at 517nm using spectrophotometer.

Methanol (2ml) with DPPH solution (0.004%, 2 ml) was used as blank. The optical density was recorded and the % inhibition was calculated using the formula given.

$$AA\% = \frac{A_b - A_s}{A_b} \times 100$$

Where AA% = Antioxidant activity (%)

A_b = Absorbance of the blank

A_s = Absorbance of the sample

Acute Toxicity

The acute toxicity study was conducted according to the method describe by Lork, 1983 [10]. The study was conducted in two phases using total of sixteen rats. In the first phases nine rats were divided in to three groups of thrice. Group 1, 2 and 3 were given 10, 100 and 1000ml/kg of body weight of the extract respectively. In the second phases 1600, 2900 and 500mg/kg were giving to the three rats respectively to determine the correct LD₅₀ values. All the treatments were given through interpretoneal. All animal were frequently observed on the day of treatment and surviving animal were further monitored daily for five days for the other sing of acute toxicity.

Statistical analysis

The experiments were done in triplicate. The result are given as mean +_ standard deviation (SD).

Result

Table 1: The mean values of antioxidant free radical scavenging of the stem of ficus iteophylla by DPPH.

Extract	Percentage (%) DPPH it scavenged (mean +standard deviation)
Acetone	89.1+0
Methanol	84.6+0.5
Ascorbic acid	91.8+0.05

Table 2: Acute lethal effect of methanol extract of *Ficus iteophylla*.

Phase one		
Doses (mg/kg)	No. of Rat	Raton of Rat dead rate
10	3	0/3
100	3	0/3
1000	3	2/3
Phase 2		
1600	1	1
2900	1	1
5000	1	1

LD₅₀ = $\sqrt{a \times b}$ (where a is the hihest non lethal doses and b is least lethal doses of the extract administerd.)

$$= \sqrt{100 \times 1000} = 316\text{mg/kg (Oral)}$$

Discussion

The uses of DPPH provided an easy and rapid way to evaluate antioxidant. The result of this investigation shows that the stem of *Ficus iteophylla* possess appreciable antioxidant activity agaist DPPH. The DPPH contains an odd electron which is responsible

for purple color and the absorbance wavelength of 517nm (C. F. R. Ferreira. *et al.*, 2007) [5]. Singh *et al.*, 2011 [14] studied the antioxidant of properties of using DPPH assay and found that different phytochemicals in leaves are responsible for higher antioxidants. Various studies have reported the presence of flavonoids, tannins, terpene and alkaloids in the extracts of *Ficus iteiphylla*. The flavonoids have been shown to possess antimicrobial, and antioxidants activities in various studies (Lin *et al.*, 2008, Lopez, 2004, Yoshidu *et al.*, 2009 and Amoral *et al.*, 2009) [18, 3]. The presence of alkaloids also had shown an antioxidant activity (Maiza *at al.*, 2007) [11] The presences of tannin in the extract may explain the antioxidant activity of the extracts as tannins are known to possess antioxidant properties (Zhang and Lin, 2008) [20], The saponins have also shown antioxidant activity (Gulcin *et al.*, 2004) The acute toxicity study of the methanolic stem extract reveal the LD₅₀ value of 316.22 mg/kg suggested by Matsumura (1975) [12] who classified chemicals base on their LD₅₀ values, and point out that LD₅₀ of 500-5000 mg/kg as slightly toxic to the experimental model. Weaknesses may be attributed to the phytochemicals constituents of the extract like tannins and saponins which are anti-nutritional factors (Umaru *et al.*, 2007).

Conclusion

The present study show that the *Ficus iteophylla* is good source of antioxidant and the results of the acute toxicity show that the plant does not cause any problem to experimental animal. Therefore the plant can be uses in traditional medicine.

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