

# Therapeutic Effects of *Viscum album* Combined with *Zingiber officinale* against CCL<sub>4</sub> induced Liver Injury in Albino Rats

## Abstract

This study was aimed at evaluating efficacy of *Viscum album* (mistletoe) and *Zingiber officinale* (ginger) in treating Rat liver against CCL<sub>4</sub>-induced liver injury. Mistletoe and ginger are used in traditional medicine for the treatment of various disorders, including hepatic disorders. Biochemical parameters and histological structure were assessed and used as a measure of therapeutic potential of the herbs against CCL<sub>4</sub>-induced liver injury. The experimental animals (15 male wistar Albino Rats) weighing between 100-120g were randomly divided into nine (3) groups. Each group comprised 5 rats and was labeled as group 1, 2 or 3. Group 1 (negative control) animals were administered saline orally daily for 6 weeks (1ml volume per kg body weight) while group 2 (CCL<sub>4</sub> group) animals were administered CCL<sub>4</sub> mixed with olive oil as vehicle in 1:1 ratio (3ml/kg body weight). Group 3 represented the treatment group with extracts of the two herbal plants (250mg/kg daily). The combined herbal extracts administered orally for 6 weeks showed a significant decrease ( $P<0.05$ ) in the concentrations or activities of liver function parameters including serum ALT, ALP, AST and GGT activities and serum protein, albumin, and bilirubin concentrations as compared with the marked increases in the parameters in CCL<sub>4</sub>-only treated rats. Histological examination of the liver of CCL<sub>4</sub>-treated rats with the combined herbal extracts showed less destruction of liver architecture in comparison to the group treated with CCL<sub>4</sub> only. The results indicated that the combined herbal extracts investigated (mistletoe, and ginger) had therapeutic effect against CCL<sub>4</sub>-induced liver injury and this effect could be due to the phytochemicals present in the herbs.

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## 1. Introduction

The liver is one of the major vital organs for the overall physiological functions of the body. It is often regarded as a “silent organ”, because it lacks inner nerve endings, it can tolerate physiological attacks and still function even at 30% of its full capacity (Wu & Cheng, 2011). Hepatotoxic agents can react with the basic cellular components and consequently induce almost all types of liver lesions. Toxins and drugs are among the basic etiopathogenetic agents of acute liver failure (Ishak & Irey, 1992). Carbon

tetrachloride (CCL<sub>4</sub>) is an occupational chemical agent widely used as a solvent in insecticide. The hepatotoxicity of halogenated hydrocarbons, particularly CCL<sub>4</sub>, has been the subject of numerous investigations in experimental animals” (Jenner & Timbrell, 1995). In ages past, nature has been a relevant source of medicinal agents and an impressive number of modern drugs have been isolated from natural sources, many based on their use in traditional medicine. A number of studies have shown that plant extracts having antioxidant activity protect against CCL<sub>4</sub> hepatotoxicity by inhibiting lipid peroxidation and enhancing antioxidant enzyme activity. Studies have shown that regular consumption of fruits, vegetables and seeds can help prevent the risk of many diseases due to their content of bioactive compounds (Peng *et al.*, 2013). The scientific evidence regarding mistletoe’s use promising. Ginger, (*Zingiber officinale* Roscoe, *Zingiberaceae*) is one of the important medicinal plant that naturally occurs in various country like India, China, South East Asia, West Indies, Mexico and other parts of the world. Ginger plants are generally 1-3 ft. in height and having different chemical constituents like Amaldehyde, Gingerol, Shogaol, and Paradol etc.

Ginger has some tremendous beneficial effect to human body to cure various types of diseases. Ginger bears an enormous number of pharmacological activities such as Neuro-protective activity and activity against colon cancer have facilitated the extent of further research for finding out less toxic and more potent drugs for the better treatment of those diseases. The seeds are also used in folk medicine, many herbal formulations and have potential therapeutic benefits due largely to the activity of their flavonoids and other bioactive compounds” (Akintonwa & Essien, 1990; Tona *et al.*, 1999; Farombi *et al.*, 2000; Pietta, 2000; Okunji *et al.*, 2002; Farombi *et al.*, 2002; Adejoke *et al.*, 2015)

**(YOUR INTRODUCTION IS NOT EXPLICIT ENOUGH, NO LITERATUIRE ON VISCUM ALBUM, AND NO OBJECTIVE OF THE STUDY)**

## **2. Materials and Methods**

### **2.1 Study Area/population**

The study was conducted at Department of Human physiology, University of Port Harcourt. It was a biological trial with Albino Wistar rats which were considered the choicest animals for this experiment because of their availability, cost, genetic makeup,

handling technique and nature of the study. Fifteen (15) healthy matured male albino wistar rats of 12 weeks old weighing between 100-120g were used in this study. The rats were obtained from the Experimental Animal Unit of the University. The rats were housed in conventional wire mesh cages under standard laboratory conditions and were allowed free access to water and feed throughout the period of the experiment

## 2.2 Consent and Ethical Consent

Formal approval was obtained from the Department of Biochemistry University of Port Harcourt for the biochemical analysis of the specimen for toxicity studies.

## 2.3 Preparation of Ethanolic Extract of *Garcinia Kola* (IS THIS GINGER?????)

### 2.4

Two kilograms (2kg) of powdery form of the *ginger* was processed at the Department of Pharmacognosy Laboratory of University of Port Harcourt for extraction using Soxhlet extraction method. During the extraction 70% of ethanol and 30% of water were used for the maceration. Two kilograms (2kg) of the seeds was macerated with ethanol and water then allowed to stand at room temperature for a period of 3 days with frequent stirring until the soluble matter dissolved. The mixture then was sieved, the damp solid material was pressed, and the solvent was clarified by filtration. The solvent was then placed in the reservoir of soxhlet for extraction. The liquid extract in the reservoir was subjected to heat for several minutes in order to vapourize the moisture. The sample was evaporated over the water bath at a temperature of 45°C and was constantly monitored until a gelatinous extract was formed.

## 2.4 Preparation of Aqueous Leaf Extract of Mistletoe (*Viscous album*)

Two kilograms (2kg) of powdery form of the **Mistletoe leaves** was taken to the Department of Pharmacognosy laboratory of University of Port Harcourt for extraction. During the extraction water was used for the maceration. Two kilograms (2kg) of the leaf was macerated with water then allowed to stand at room temperature for a period of 3 days with frequent stirring until the soluble matter dissolved. The mixture then was sieved, the damp solid material was pressed, and the solvent was clarified by filtration.

The extract was then placed in the reservoir of soxhlet for extraction. The liquid extract in the reservoir was subjected to heat for several minutes in order to vapourize the moisture. The sample was evaporated over the water bath at a temperature of 45<sup>0</sup>C and was constantly monitored until a gelatinous extract was formed.

## **2.5 Grouping and Treatment of Animals**

Fifteen (15) male Albino Wistar Rats were used for this research and were divided according to their body weight into 3 groups with each group containing five (5) Rats.

**Group 1:** This was the negative control group; they received 1ml of distilled water daily for six (6) weeks.

**Group 2:** This group was induced with Carbon tetrachloride (CCL<sub>4</sub>) causing hepatotoxicity using 3ml/kg body weight and served as a positive Control.

**Group 3:** 24 hours after inducing with CCL<sub>4</sub> this group received 250mg/kg body weight each of both Mistletoe and *Garcinia Kola* and daily for six (6) weeks.

In the studied animals, hepatic injury in all groups except standard control was induced by single oral administration of CCL<sub>4</sub> mixed with olive oil as vehicle in 1:1 ratio (3 ml/kg of rat body weight. A pilot study was first carried out using 10 Albino Wistar Rats for each of the three (3) herbs extracts used (Mistletoe, Bitter Kola). The results obtained showed that the lethal dose was estimated at 1,500mg/kg.

## **2.6 Procedures for Administration of Extracts**

Administration of extract was by oral gavage route. The rat was held at the skin over the head and turned so that the mouth was faced upward and the body lowered towards the holder. The syringe needle knob was then placed into the mouth of the rat a bit laterally to avoid the teeth which are centrally located. The syringe content was then gradually emptied drop by drop into the mouth of the rat.

## **2.7 Sample Collection**

The blood samples were collected from the animal via cardiac puncture and sacrificed under 70% chloroform anesthesia into plain specimen bottle. The samples were allowed to clot, then centrifuged at 3000 revolutions per minute for 3 minutes. Then sera obtained were stored in a freezer until required for use for analysis for liver function.

## **2.8 Histological Studies**

After 24 hours of induction and after 6 weeks of the experiment (for both controls and treatment), an animal in each group was dissected and their liver tissues were histologically studied.

## **2.9 Statistical Analysis**

The data was evaluated statistically by SPSS version 20. Using one way analysis of variance (one way ANOVA) and subjected to Fischer LSD post Hoc. Results were expressed as mean  $\pm$ SD. Difference between means were considered significant at  $P < 0.05$ .

## **3. Results**

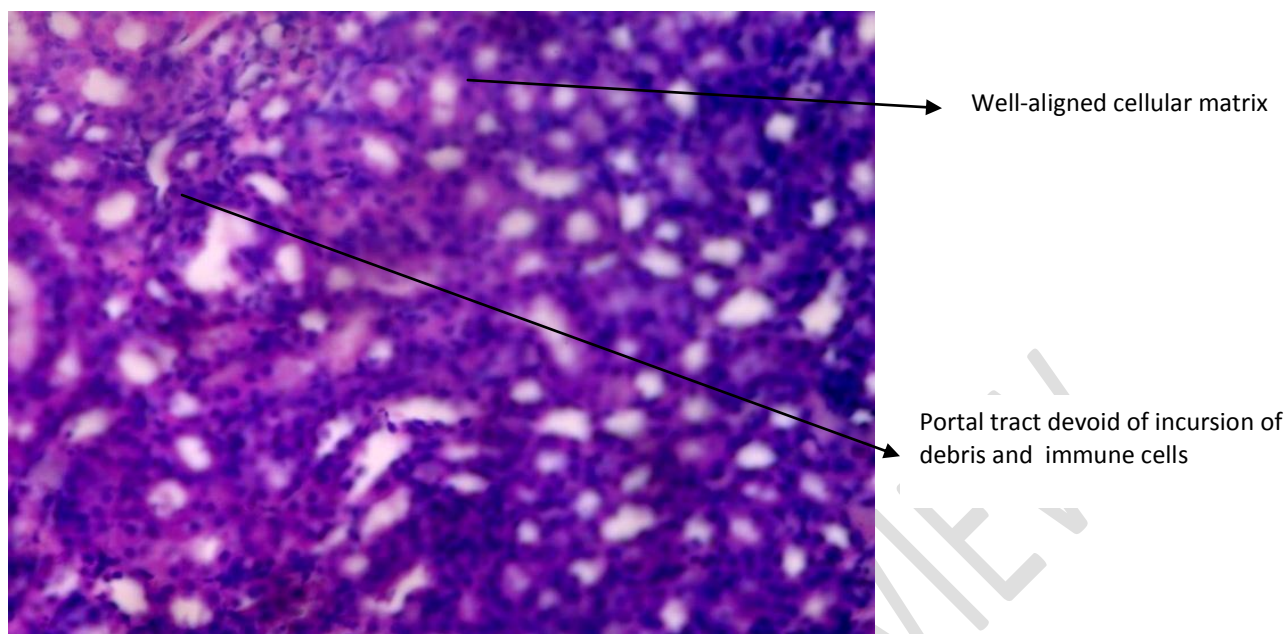
### **4.1: Comparison of Parameters for Rats Treated with Combination Extracts of Mistletoe and Ginger (Group 3) with Negative and Positive Controls.**

The Table below showed ANOVA results and Post hoc results. ANOVA results showed significant difference in the means of the three groups (Group 1, Group 2, and Group 3) while Post hoc result showed significant difference between in the means of the groups being compared except between Group 1 and Group 3 for AST.

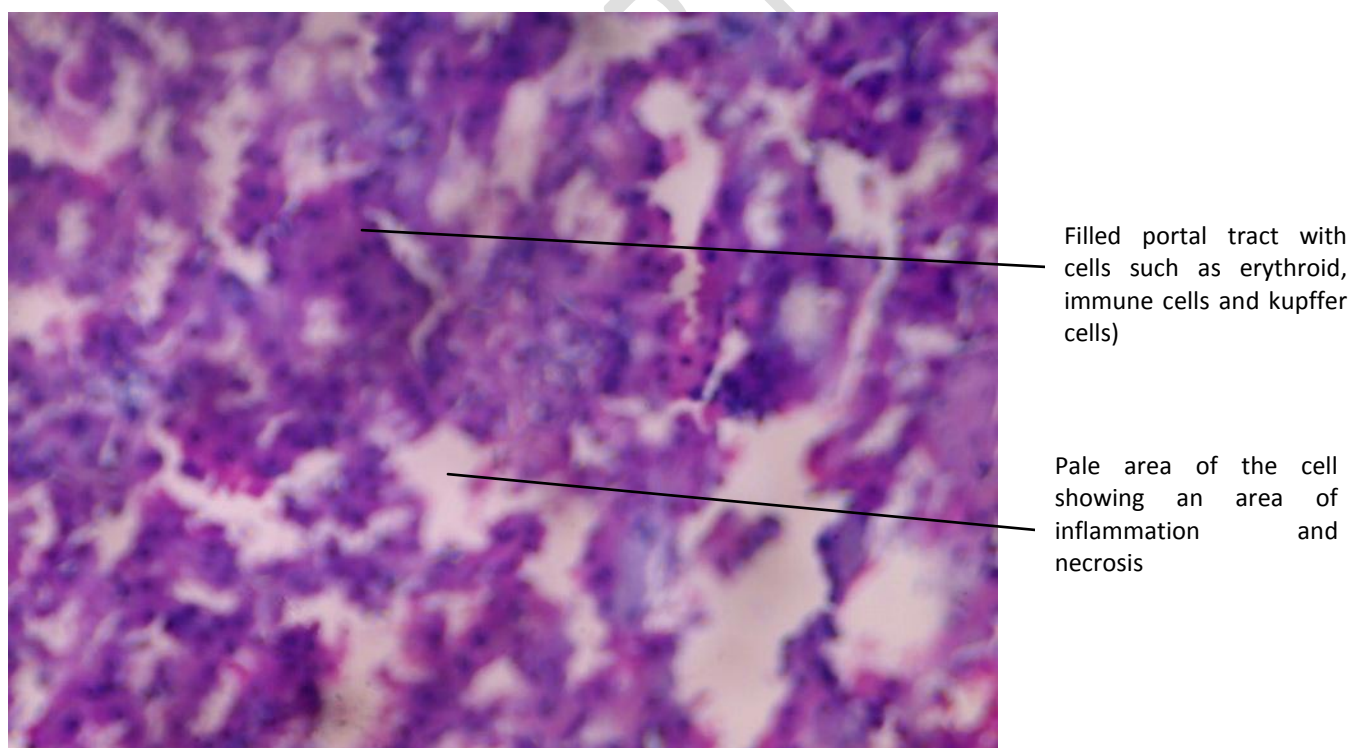
136 **Table 1: Comparison of Parameters for Rats Treated with Extracts of Mistletoe and Ginger**

| GROUPS  | Protein<br>(g/L) | Albumin<br>(g/L) | Total bilirubin<br>(mmol/L) | Conj. Bilirubin<br>(mmol/L) | ALT<br>(u/l) | ALP<br>(u/l) | GGT<br>(u/l) | AST<br>(u/L) |
|---|------------------|------------------|-----------------------------|-----------------------------|--------------|--------------|--------------|--------------|
| Group 1(Negative control)                       | 30.46± 5.44      | 25.96±5.01       | 10.41±6.09                  | 6.78±0.64                   | 7.60±0.89    | 86.0±1.00    | 27.98±0.78   | 28.40±11.50  |
| Group 2 (positive control)                      | 81.34±10.08      | 78.08±3.94       | 91.23±1.42                  | 48.99±1.95                  | 19.00±2.12   | 387.80±4.82  | 88.0±4.69    | 144.40±18.62 |
| GROUP 3 <b>mistletoe+ginger+CCL<sub>4</sub></b> | 60.60±3.83       | 32.40±1.37       | 25.30±3.75                  | 12.16±2.10                  | 11.00±0.00   | 260.6±83.62  | 36.2±1.64    | 26.40±8.29   |
| P-Value   | <0.0001          | <0.0001          | <0.0001                     | <0.0003                     | <0.0001      | <0.0001      | <0.0001      | <0.0006      |
| F-Values  | 86.92            | 18.35            | 95.67                       | 27.29                       | 30.24        | 19.11        | 28.84        | 15.23        |
| Post Hoc  | S                |                  |                             | S                           |              |              |              | S            |
| Group 1 vs Group 2                              | S                | S                | S                           |                             | S            | S            | S            | S            |
| Group 1 vs Group 3                              | S                | S                | S                           | S                           | S            | S            | S            | NS           |
| Group 2 vs Group 3                              | S                | S                | S                           | S                           | S            | S            | S            | S            |

137 Values are presented in mean ±SD, n=5 per group, S=significant difference when compared **WHERE ARE THE SUPERSCRIPTS**  
 138 **ON YOUR MEAN VALUES TO SHOW THE HIGHEST AND THE LOWEST MEANS AND THEIR SIMILARITIES**

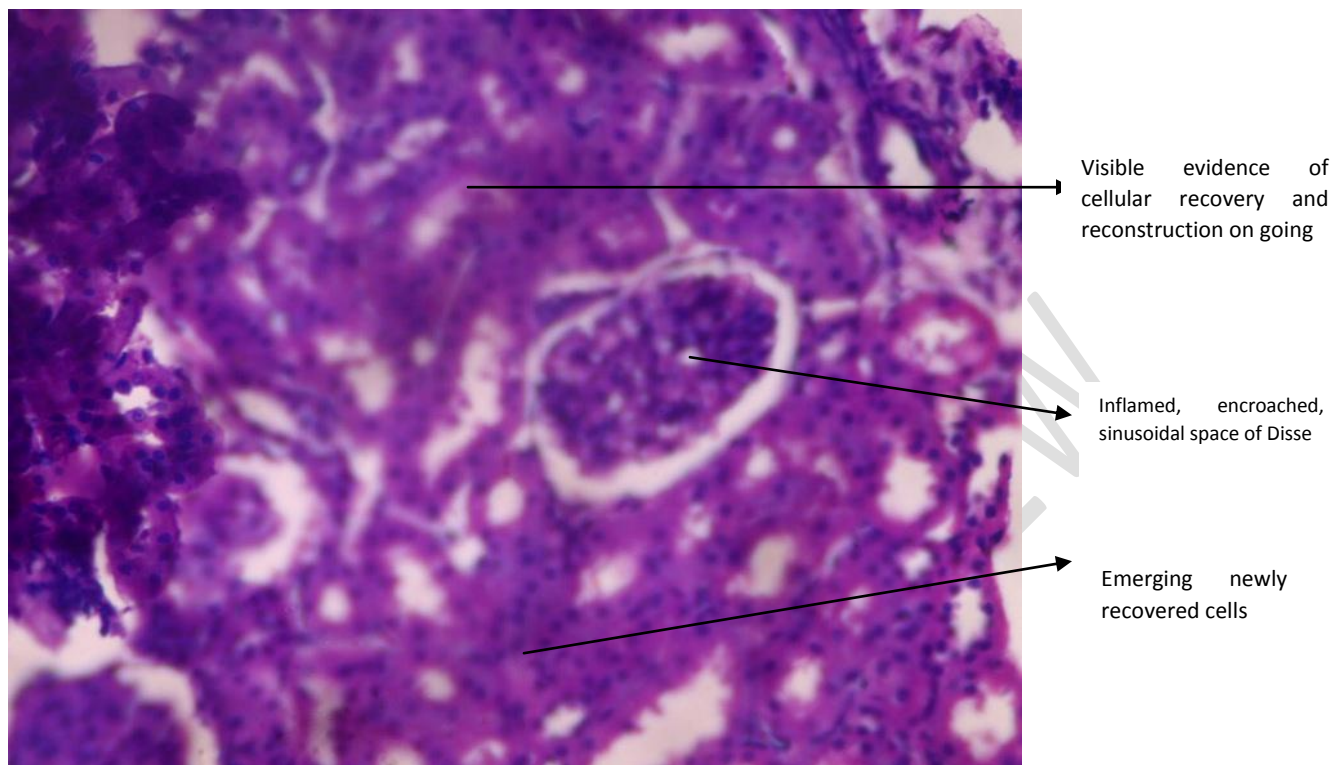


**Fig 1.** Photo micrographic slide of liver organ of group 1 (negative control saline) H & E X400



**Fig 2.** Photo micrographic slide of liver organ of group 2 (*positive control CCL<sub>4</sub>-induced hepatotoxicity*) H & E X400





**Fig 3.** Photo micrographic slide of treated liver organ of group 3 using extracts of *mistletoe*, and *ginger* (250mg/kg). H & E X400.



#### 4. Discussion

The therapeutic effects of the herbs under study; (*Viscum album* (Mistletoe) and *Zingiber officinale* (Ginger) aptly demonstrated the ability to lower the levels of alanine aminotransferase (ALT), aspartate aminotransferase (AST), and alkaline phosphatase (ALP) in Albino Wistar Rats earlier induced with CCL<sub>4</sub>. ALT, AST and ALP are parameters that could be used to monitor the liver damage and to what extent. Hence, drop in their levels are usually seen as signal of recovery. The combination of the herb extracts was observed to show potency by significantly reducing the levels of the liver enzymes (ALT, AST, ALP and GGT). The findings in this present study is similar to that reported by Edward (2006) indicated the phytochemical components of these herbs may be seen to have effective impact on hepatic recovery probably through mopping up the radicals of CCL<sub>4</sub> that are causing the damage. This interaction may give the liver the opportunity to recover. This shows that the consumption of these herbal extracts by the rats may be increasing the rate of protein synthesis leading to the higher concentration of albumin and protein in the liver.

The livers of saline control Albino Wistar Rats revealed the normal characteristic pattern of hepatic architecture (Fig 1). The liver of Albino Wistar Rats subjected to CCL<sub>4</sub> showed disrupted or loss of liver tissue architecture, severe dilatation and congestion of blood vessels (either central veins or portal tract vessels), marked lymphocytic infiltration, and fibrosis extending between the portal areas (Fig 2). Despite the incidence of necrosis inflicted by the CCL<sub>4</sub> treatment, there appeared significant evidence of cellular reconstruction and recovery, after treatment with the combination herbal extracts and consequently, there was a visible pattern of new cells emerging (Fig 3).

**(YOUR DISCUSSION IS TOO SHORT COMPARE TO THE ARRAY OF RESULT THAT YOU HAVE)**

#### 5. Conclusion

The results of the present study suggest that mistletoe and ginger preparation may be a useful therapeutic intervention for patients with acute liver disease that have similar mechanism of damage induced by CCL<sub>4</sub>. The mechanism(s) by which *Viscum album* and *Zingiber officinale* modulate hepatic inflammation remains, however, unclear, therefore, studies are required to elucidate the mechanism(s) by which mistletoe and ginger preparations exert their therapeutic

potential as seen in the present study. **(YOUR CONCLUSION IS NOT CONCISE PLEASE  
REDUCE IT TO JUST ONE SENTENCE)**

## References

- Adejoke, E. M., Ibukun, D. A., Erommonselle, E. & Adam, M. A. (2015). Hepatoprotective Effects of *Garcinia kola* (bitter kola) against Paracetamol-induced Oxidative Damage and Glycogen Degranulation in Hepatocytes of Adult Male Wistar Rats. *Journal of Advances in Biology and Biotechnology*, 3(3), 110-116.
- Akintowa, A. & Essien, A. R. (1990). Protective Effects of *Garicinia kola* Seed Extract against Paracetamol-induced Hepatotoxicity in Rats. *Journal of Enthnopharmacology*, 29 (2), 207-211.
- Edward, L. K. (2006). Schiff's diseases of the liver, 11<sup>th</sup> edition, *New England Journal of Medicine* 354 (1), 54-66.
- Farombi, E. O. (2000). Mechanisms for the Hepato Protective Action of Kolaviron Studies on Hepatic Enzymes, Microsomal Lipids and Lipid per Oxidation in Carbon Tetrachloride Treated Rats. *Pharmacological Research*, 42, 75-80.
- Farombi, E. O., Okanni, O. O. & Emerole, G. O. (2002). Anti Oxidative and Sweetening Activities of Kolaviron in Vitro. *Pharmaceutical Biology*, 40,107-116.
- Ishak, K. & Irey, N. (1972). Hepatic Injury Associated with the Phenothiazines. *Archives of Pathology*, 93 (4), 283-304.
- Jenner, A. M. & Timbrell J. A. (1995). In vitro Microsomal Metabolism of Hydrazine. *Xenobiotica*, 25, 599-609.
- Okunji, C. O., Tantalia, A. W., Hicks, R. P, Iwu, M. M. & Skanchoy, D. J. (2002). Capillary Electrophoresis Determination of Medicinal Formulations. *Plant Medicine*, 68, 440-444.
- Pietta, P. G. (2000). Flavonoids as Antioxidant. *Journal of Natural Products*, 63 (7), 1035-1042.
- Peng, W., Wu, J., Wan, Z., Yi, J. & Wu, Y. (2013). Investigation of the Extracts from *Bidens pilosa* Linn. var. *radiata* Sch. Bip for Antioxidant Activities and Cytotoxicity against Human Tumour Cells. 67 (1), 17-26.
- Tona, L., Ngmibi, N. P., Tasakala, M., Mesiak, K., Cimanga K. & Apers, S. (1999). Antimalarial Activity of Extract of 20 Crude Extracts from Nine African Medicinal Plants used in Kinshasa, Congo. *Journal of Ethnopharmacology*, 15(68), 193-203.

231 Wu, M. & Cheng, C. (2011). Liver Disease in Traditional Chinese Medicine. *709 research*  
232 *Report of Toronto School of Traditional Chinese Medicine*, 1-35.

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