

## Phytochemicals, bioactivities of *Ximenia Americana*: A review

Khaled Rashed\*

Department of Pharmacognosy, National Research Centre, 33 El-Bohouth st.-Dokki, Giza, P.O.12622, Egypt

### Abstract

The genus *Ximenia* belongs to the Olacaceae and comprises about 8 species. *Ximenia americana* is one of the most valuable wild edible plants in the world. In different countries, it is utilized as food, medicine, an essential oil source, and the industrial component to other products. It is a plant that is used in traditional medicine for the treatment of malaria, leprotic ulcers and skin infections of mixed origin in Northern parts of Nigeria. *Ximenia Americana* is reputable for the treatment of various ailments in Nigeria. It has been alleged to have antineoplastic activities, antimicrobial, anticancer anti-inflammatory activities and lots more.

**Keywords:** *Ximenia americana*, chemical compounds, bioactivities.

#### \*Correspondence Info:

Dr. Khaled Rashed  
Department of Pharmacognosy,  
National Research Centre, 33 El-Bohouth st.-  
Dokki, Giza, P.O.12622, Egypt

#### \*Article History:

**Received:** 29/10/2021  
**Revised:** 27/01/2022  
**Accepted:** 26/02/2022  
**DOI:** <https://doi.org/10.7439/ijbar.v13i4.5701>

#### QR Code



**How to cite:** Rashed K. Phytochemicals, bioactivities of *Ximenia Americana*: A review. *International Journal of Biomedical and Advance Research* 2022; 13(04): e5701. Doi: 10.7439/ijbar.v13i4.5701 Available from: <https://ssjournals.com/index.php/ijbar/article/view/5701>

Copyright (c) 2022 International Journal of Biomedical and Advance Research. This work is licensed under a [Creative Commons Attribution 4.0 International License](https://creativecommons.org/licenses/by/4.0/)

### 1. Introduction

*Ximenia americana* belongs to the family Olacaceae. The genus *Ximenia* has eight different species namely *Ximenia roigi*, *Ximenia aegyptiaca*, *Ximenia parviflora*, *Ximenia coriacea*, *Ximenia aculeate*, *Ximenia caffra*, *Ximenia americana* and *Ximenia aegyptica* [1]. It is commonly referred to as wild plum and tallow nut. It is a shrub or small tree that grows to about 7 m high, with zigzag branches. The bark is black or grey brown, smooth when young but becomes rough with age having stiff axillary spines. The leaves are alternate or clustered on spur shoots. The seed morphology is variable. Flowers are green white to greenish yellow, scented and 5-10 mm long in small branched inflorescences [2].

Native names in Nigeria: Hausa (Tsada), Fulani (Chabull), Tiv (Anomadze), Yoruba (Igo) and Benin (Alimo-mamiwota) [3]. The methanolic extract of leaves of *Ximenia americana* has been proven to have anti-diabetic effect in rats [4]. The ethanolic extract of the bark of *X. americana* revealed the presence of the following metabolites; flavonoids, steroids, tannins, alkaloids, phenolic compounds, saponin, terpenoids, and glycosides. It also has activity against *Staphylococcus aureus* and low

activity against *Pseudomonas aeruginosa*. Other investigations led to isolation of two compounds. The first one which is new is 3-methyl-1-oxoisochroman-8-carboxylic acid and the second compound which is a known steroid is ergosta-4,6,8,22-tetraen-3-one [5]. The plant has also been reported to have anti-HIV/AIDS related diseases effects such as abscesses, skin rashes, diarrhea and gonorrhea. The anti-HIV/AIDS effect of *Ximenia americana* is due to the fact that it contains oleic, ximenic (hexacos-17-enoic), linoleic, linoleic and stearic acids. Its oil consists of very long chain fatty acids with up to 40 carbon atoms used to manage STIs including gonorrhoea [6].

This review article is aimed at addressing the phytochemicals and bioactivities of *Ximenia Americana*. It has been alleged to have antineoplastic activities, antimicrobial and anti-inflammatory activities and lots more. The aqueous plant extract of the leaves is used in the treatment of cancer in African traditional medicine [7]. *X. Americana* has a wild varieties of phytochemical compounds which among them are the following secondary metabolites; saponins, flavonoids, tannins, terpenoids,

sterols, quinines, alkaloids, cyanogenetics glycosides, cardiac glycosides and carbohydrates in the form of sugars and soluble starch [8]. The essential oils of the leaves of *X. americana* using GC-MS and identified 33 components representing 98% of the total oil. The main constituents analysed are benzaldehyde 63.5%, hydroxybenzylcyanide 13%, isophorone 3.5%. The hydroxybenzylcyanide is known as a primary breakdown products of glucosinolates found mainly in the Brassicaceae family [9]. The fixed oil present in the seeds of *X. americana* [10].

## 2. Chemical compounds and Bioactivities

The stem ethanolic extract of the plant afforded steroids (stigmasterol and sitosterol), triterpenoids (betulinic acid, oleanolic acid, 28-O-( $\beta$ -D-glucopyranosyl) oleanolic acid, 3-oxo-oleanolic acid, 3 $\beta$ -hydroxycycloart-24(E)-ene-26-oic acid and sesquiterpenoids (furanic and widdrane type). A large number of sesquiterpenes are constituents of essential oils of higher plants and seem to intervene in the pharmacological properties attributed to these volatile fractions [11]. It has been clarified that the biological activities of the liverworts are due to terpenoids and lipophilic aromatic compounds [12]. Steroids and triterpenes with therapeutic interest and manufacturing employment, are a group of secondary metabolites of outstanding importance [11]. Considerable recent work strongly indicates the great potential of the triterpenoids as source of use medicinal [13].

In general, the compounds found in *X. americana* were saponins, glycosides, flavonoids, tannins, phenolics, alkaloids, quinones and terpenoids types. In addition, the plant is potentially rich in fatty acids and glycerides and the seeds contain derivatives cyanide. The identified compounds did not demonstrate a representative pattern of each class. For example, the sesquiterpene were furanolic and widdrane while, triterpenes exhibited oleanane and cycloartane skeletal type. Concerning the fatty acids, in addition to common C<sub>16</sub>, C<sub>18</sub> and C<sub>22</sub>, a distinctive feature is the presence of acetylenic, as well as, very long chain fatty acids.

### 2.1 Antimicrobial and antifungal potentials

The MeOH extract from leaves of *X. americana* inhibited or retarded growth of *Neisseria gonorrhoea* organism at dilution as low as 250  $\mu$ g/ml. This same extract showed antifungal effect against *Candida albicans* and *Cryptococcus neoformans* in concentration of 4000  $\mu$ g/ml. Chemical screening conducted on the extract showed the presence of several secondary metabolites as tannins, sterols, terpenoids, flavonoids and saponins [14].

### 2.2 Analgesic activity

The aqueous extract of stem bark of *X. americana* has analgesic properties that justify its use popular in

countries such as Tanzania, Senegal, Zimbabwe and Nigeria. The extract of *X. americana* in doses containing 10 to 100 mg/kg P.C, inhibits contractions of the abdomen with analgesic effects comparable to those of phenylbutazone. In fact, at doses of 100 mg / kg P.C, phenylbutazone causes an inhibition of pain in 45.2 $\pm$ 2%. The percentage of inhibition by extract of *X. americana* is 61.1 $\pm$ % in the same concentration. These properties are probably due to the presence of flavonoids and saponins, detected in the extract.

The analgesic activity of the methanol extract of *X. americana* leaf was investigated in chemical models of nociception in mice. The extract at doses of 200, 400 and 600 mg/kg i.p. produced an inhibition of 54.13, 63.74, and 66.4% respectively, of the abdominal writhes induced by acetic acid in mice. In the formalin test, the administration of 200, 400 and 600 mg/kg i.p. had no effects in the first phase (0 to 5 min) but produced a dose dependent analgesic effect on the second phase (15 to 40 min) with inhibitions of the licking time of 29.3, 47.8 and 59.8%, respectively. These observations suggested that methanol extract of *X. americana* leaf possesses analgesic activity [15].

### 2.3 Antipyretic activity

The bark of stem of *X. americana* has been used in West Africa for the treatment of pain and fever. To verify this second property, the treatment of rats in hyperthermia with *Ximenia americana* stem bark aqueous and with beer yeast was compared to those obtained with lysine acetylsalicylate (Aspegic). The study showed an antipyretic action of the extract. Moreover, the toxicological study of the stem extract indicated a LD<sub>50</sub> of 237.5 mg/kg P.C according to the classification of Diezi this plant is relatively toxic. The experiments show that the properties of *X. americana* could be due to the presence of saponins, as shown by screening tests performed in this study. These results justified the use of *X. americana* in traditional cure of fever treatment [16].

### 2.4 Antitrypanosomal potential

The *in vitro* antitrypanosomal activity of methanolic and aqueous extracts of stem bark of *Ximenia americana* was evaluated on *Trypanosoma congolense*. Blood obtained from high infected mice with *T. congolense* was incubated with methanolic and aqueous extracts at 20, 10 and 5 mg/ml and Dimininal(R) (diminazene aceturate) at 200, 100 and 50  $\mu$ g/ml in a 96 micro plate. The results revealed that methanol and aqueous extracts had activity at 20 and 40 mg/ml however, the methanolic extracts were more active than aqueous extracts at 10 and 5 mg/ml. Phytochemical screening of the methanolic and aqueous extracts of the bark showed that they both had flavonoids, anthraquinones, saponins, terpenes and tannins. The aqueous and methanolic extract appears to show some potential activity against *T. congolense* [17].

## 2.5 Anticancer activity

Two sesquiterpenes isolated from the EtOH extract of the stems of *X. americana* did not inhibit the growth of HL-60 (human leukemia), HTC-8 (human colon) and MDA-MB-435 (human breast cancer) cell lines. The antineoplastic activity in vitro of various extracts from *Ximenia americana*, plant used in African traditional medicine for the treating cancer, was investigated [18]. The most active, aqueous extract was subjected to a detailed investigation in a panel of 17 tumor cell lines originating from human (16 lines) and rat (1 line), showing an average IC<sub>50</sub> of 49 mg raw powder/ml medium. The majority of cell lines (11 out of 17) were classified as sensitive (the sensitivity varied from 1.7 mg/ml in MCF7 breast cancer cells to 170 mg/ml in AR230 chronic-myeloid leukemia cells) and three of these (MCF7 breast cancer, BV173 CML and CC531 rat colon carcinoma) showed a particularly high sensitivity, with ratios lower than 0.1 of the average IC<sub>50</sub>. The *in vivo* antitumor activity was determined in the CC531 colorectal rat model and significant anticancer activity was found following per oral administration, indicating a 95% reduced activity [18].

## 2.6 Antiviral effect

The stem bark MeOH extract of *X. americana* as well as several others plant species used by the Maasai pastoralis of East Africa showed antiviral effect against measles virus in vitro by plaque reduction neutralization assay. Potentially active constituents from extracts of all the plants include polyphenols, alkaloids, tannins, sterols, terpenes, saponins and glycosides, between others [19].

## 2.7 Hepatic and hematological effects

A study was conducted from the leaves, stem bark and root aqueous extract of *X. americana* with albino rats. The results of this work shows that the extracts significantly ( $P < 0.05$ ) increasing the level of serum alanine transaminase (ALT) and aspartate transaminase (AST), results indicative of hepatocellular damage. The result also shows that the root has the ability to impair albumin synthesis as observed by the decrease of level of serum albumin. The weight of the animal showed a significant ( $P < 0.05$ ) reduction on administering the leaves extract as compared to the control and the others extracts. This reduction might be due to poor intake and utilization of food by the animals in the leaves extract group. The significantly ( $P < 0.05$ ) higher content of hydrogen cyanide, saponins, and oxalates in the root extracts indicates that the root extracts may be more toxic. Hydrogen cyanide is known to cause gastrointestinal inflammation and inhibition of cellular respiration. Saponins are known to have haemolytic properties and the ability to reduce body cholesterol by preventing its reabsorption. The high saponin content in the root may lead to gastroenteritis manifested by diarrhea. Oxalates have

been known to cause irreversible oxalate nephrosis when ingested in large doses. Thus, there is need to isolate the specific component(s) responsible for the toxicity in the root extract in order to standardized the preparation for maximum therapeutic benefit [20].

## 2.8 Food composition and cosmetic use

Glyceride blends containing ximenynic acid (found in *X. americana*) are useful for the preparation of food compositions or food supplements, including margarine, chocolate, ice cream, mayonnaises, cheese, dry soups, drinks, cereal bars and sauces and snack bars. The blend provides a composition providing health benefits consisting of insulin resistance, or related disorders such as diabetes, delaying the onset of symptoms related to development of Alzheimer's disease, improving memory function, lowering blood lipid levels, anticancer effects or skin antiageing effects. Foods *X. americana* flowers are a replacement for orange blossoms with similar fragrance and soothing cosmetic properties [21].

## 3. Conclusion

*Ximenia americana* has several medicinal uses as anticancer, antioxidant, antimicrobial and others. The plant could be a veritable and cheaper substitute for conventional drugs since the plant is easily obtainable and the extract can easily be made via a simple process of maceration or infusion.

## References

- [1]. Irvin FR. Woody plant of Ghana. London Oxford University Press, 1961: 472-474.
- [2]. Ehigiamusoe RE. A herbarium of Nigeria medicinal plants. University of Calabar Press, 2013: 478.
- [3]. Siddaiah M, Jayaveera KN, Souris K, Yashodha Krishna JP, Vasanth Kumar P. Phytochemical screening and anti diabetic activity of methanolic extract of leaves of *Ximenia americana* in rats. *Int J Innov Pharmac Res.* 2011; 2(1): 78-83.
- [4]. Abdalla AA, Shyaula SL, Ishak CY, Ayoub SMH. Bioassay and phytochemical studies on *Ximenia americana* L. Bark ethanolic extract. *J Forest Prod Indust.* 2013; 2(3): 63-68.
- [5]. Chinsembu KC. Ethnobotanical study of plants used in the management of HIV/AIDS-related diseases in Livingstone, Southern Province, Zambia. *Evid-Based Complem Altern Med.* 2016: 4238625.
- [6]. James DB, Owolabi AO, Ibiyeye H, Magaji J, Ikugiyi YA. Assessment of hepatic effect, hematological and some phytochemical constitute of *X. americana* (leaves, stem and root extract). *Afr J Biotechnol.* 2008; 23: 4274-4278.

- [7]. Ogunleye DS, Abitoye D. Studies of antimicrobial activities and chemical constitute of *X. americana*. *Trop J Pharmac Res*. 2003; 2(2): 239-241.
- [8]. Geyid A, Abebe D, Debela A, Makonnen Z, Aberra F, Biza T, et al. Screening of medicinal plants of Ethiopia for their anti-microbial properties and chemical profiles. *J Ethnopharmacol*. 2005; 97(3): 421-427.
- [9]. Mevy J-P, Bessiere J-M, Greff S, Zombre G, Viaria J. Composition of the volatile oil from the leaves of *Ximenia americana* Linn. *Biochem Syst Ecol*. 2006; 34: 549-553.
- [10]. Saeed AEM, Bashier RSM. Physico-chemical analyses of *Ximenia americana* Linn, seed oil and structure elucidation of some chemicals constituents of its seed oil and fruit pulp. *J Pharmacognosy Phytother*. 2010; 2(4): 49-55.
- [11]. Bruneton, J. (3a Ed) (1999). Pharmacognosie, phytochimie, plantes médicinales, Tec & Doc Ed., Angers, France.
- [12]. Atta-ur-Rahman. (1988). Studies in Natural Products Chemistry, Structure Elucidation, Vol.32, Elsevier, New York, U.S.A.
- [13]. Mahato, S. B.; Nandy, A. K. & Roy, G. Triterpenoids. *Phytochemistry*, 1992; 31: 2199-2249,
- [14]. Geyid, A.; Abebe, D.; Debella, A.; Makonnen, Z.; Aberra, F.; Teka, F.; Kebede, T.; Urga, K.; Yersaw, K.; Biza, T.; Mariam, B. H. & Guta, M. Screening of medicinal plants of Ethiopia for their anti-microbial properties and chemical profiles. *Journal of Ethnopharmacology*, 2005; 97: 421-427.
- [15]. Siddaiah, M.; Jayavcera, K. N.; Mallikarjuna, R. P.; Ravindra, R. K.; Yasodha, K. Y. & Narender, R. G. Phytochemical screening and analgesic activity of methanolic extract of *Ximenia americana*. *Journal of Pharmacy and Chemistry*, 2009; 3(1): 23-25.
- [16]. Soro, T. Y.; Traore, F.; Datte, J. Y. & Nene-Bi, A. S. Antipyretic activity of aqueous extract of *Ximenia americana*. *Phytoterapie*, 2009; 7 (6): 297-303.
- [17]. Maikai, V. A.; Kobo, P. I. & Auda, A. O. Acute toxicity studies of aqueous stem bark extract of *Ximenia Americana*. *African Journal of Biotechnology*, 2008; 7 (10): 1600-1603.
- [18]. Voss, C.; Eyol, E. & Berger, M. R. Identification of potent anticancer activity in *Ximenia americana* aqueous extracts used by African traditional medicine. *Toxicology and Applied Pharmacology*, 2006; 211: 177-178.
- [19]. Parker, M. E.; Chabot, S.; Ward, B. J. & Johns, T. Traditional dietary additives of the Maasai are antiviral against the measles virus. *Journal of Ethnopharmacology*, 2007; 114: 146-152.
- [20]. James, D. B.; Owolabi, A. O.; Ibiyeye, H.; Magaji, J. & Ikugiyi, Y. A. Assessment of the hepatic effects , hematological effect and some phytochemical constituents of *Ximenia Americana* (Leaves, stem and root) extracts. *African Journal of Biotechnology*, 2008; 7: 23: 4274-4278.
- [21]. Paolo, R. Cosmetic use of the oil and flowers of *Ximenia americana*. *Rivista Italiana Essenze*, 1979; 61; 5: 190-193.