

*Received*: Sept, 2024 *Accepted for Publication*: Dec. 2024

# DPPH Radical Scavenging Activities Of The Air-dried Leaves Essential Oil From *Taminalia catappa* LINNAEUS and *Alternanthera pungens* KUNTH

\* <u>Ogunmoye</u> A. O.<sup>1</sup>, <u>Olubomehin</u> O. O.<sup>1</sup>, <u>Atewolara-Odule</u> O.C.<sup>1</sup>, <u>Ibikunle</u> A. A.<sup>1</sup>, <u>Sanyaolu</u> N. O., <u>Yussuf</u> S. T.<sup>1</sup>, <u>Onajobi</u> I. B.<sup>2</sup> and Rowaiye G.<sup>3</sup>

 <sup>1</sup>Department of Chemical Sciences, Olabisi Onabanjo University, P.M.B. 2002, Ago-Iwoye, Ogun State, Nigeria.
<sup>2</sup>Department of Microbiology, Olabisi Onabanjo University, P.M.B. 2002, Ago-Iwoye, Ogun State, Nigeria.
<sup>3</sup>National Biotechnology Research Development Agency, Federal Ministry of Innovation, Science and Technology, Nigeria
\*Author for Correspondence: omotunde.ogunmoye@oouagoiwoye.edu.ng

## ABSTRACT

Essential oils are aromatic, volatile, and lipophilic compounds extracted from various parts of plants, such as flowers, leaves, stems, roots, seeds, bark, and fruits. They are highly valued for their flavor, therapeutic, and aromatic properties, and are widely used in food, medicine, and cosmetic industries. In this study, essential oils were extracted from the leaves of *Terminalia catappa* and Alternanthera pungens using hydrodistillation with an all-glass Clevenger-type apparatus. The oils were then assessed for antioxidant activity using the 2,2-diphenyl-1-picrylhydrazyl (DPPH) assay, with ascorbic acid as the reference standard. The extracted oils appeared as cloudy light yellow to cloudy white liquids with a strong aroma. Yield percentages were 0.25% for *T. catappa* and 0.40% for A. pungens. The inhibition percentages ranged from 84.67% at 25 mg/mL to 54.22% at 100 mg/mL for *T. catappa*, and from 73.87% at 100 mg/mL to 43.90% at 50 mg/mL for A. pungens. The antioxidant evaluation revealed moderate to high radical scavenging activity compared to the standard. Overall, the results highlight the potential of these essential oils as natural antioxidants suitable for applications in food, pharmaceutical, and cosmetic products.

Keywords: Medicinal plant, Essential oil, DPPH, Antioxidant Activity

# **INTRODUCTION**

Essential oils, also known as volatile aromatic oils, are fragrant oily substances extracted from various parts of plants. The extraction process is often labor-intensive and time-consuming, and the method used is crucial, as certain solvent-based techniques may compromise the oils' therapeutic properties. The chemical composition of essential oils varies significantly, with major constituents making up as much as 85% of the oil, while others are found only in trace amounts (Miguel, 2010).

Antioxidants are compounds that neutralize free radicals by donating electrons, transforming them into harmless molecules. A growing body of evidence suggests that cellular damage caused by free radicals contributes to the development of various conditions, including neurological disorders, cancer, cardiovascular diseases, and weakened immune function (Aruoma, 1998; Kamatou and Viljoen, 2010). Oxidative stress, in particular, plays a key role in chronic and degenerative diseases such as cancer, autoimmune conditions, rheumatoid arthritis, cataracts, aging, and neurodegenerative disorders (Willcox et al., 2004; Pham-Huy et al., 2008).

The antioxidant capacity of phenolic compounds is primarily attributed to their redox behavior, enabling them to act as reducing agents, hydrogen donors, singlet oxygen quenchers, and metal ion chelators (Kumar, 2005; Isah, 2021). As a result, naturally occurring antioxidants have gained considerable attention for their potential to protect the human body from oxidative damage and for their ability to safeguard food products against deterioration caused by oxidation (Osawa et al., 1990; Houghton, 1995; Maestri et al., 2006).

Terminalia catappabelongs to the Combretaceae family and is a perennial tree species that thrives in tropical climates, making it widespread across nearly all regions of the country. Terminalia catappa is also known by several other names, including tropical almond, wild almond, Indian almond, sea almond, beach almond, and Malabar almond (Untwal and Kondawar, 2006; Orwa et al., 2009). The plant is refers to as "Afonzon" in Yoruba land, Southwest Nigeria(Gbile, 1984; Burkill, 1985). The juice from the plant is used in the preparation of ointment for scabies, leprosy and other cutaneous diseases (Nair and Chanda, 2008). Terminalia catappa has been reported to possess various medicinal properties, including antimicrobial, antidiabetic, antinociceptive, antiparasitic, antifungal, and antibacterial activities (Elizabeth, 2005; Rao and Nammi, 2006; Rajarajan et al., 2010). It also exhibits antioxidant effects (Ko et al., 2002; Mety and Mathad, 2011), anticancer potential (Chu et al., 2007), and is used in the treatment of hepatitis and other liver-related ailments in Taiwan (Lin and Kan, 1990). Additionally, the tree is commonly cultivated in southeastern Nigeria for shade and ornamental purposes (Ezeokonkwo and Dodson, 2004; Agu and Menkiti, 2017).

Various phytochemicals have been identified in extracts from the leaves of Terminalia catappa, including gallic acid, triterpenic acids, 4hydroxyphenylpropionic acid, m-coumaric acid, 3,4-dihydroxybenzoic acid, p-coumaric acid, kaempferol, quercetin, tergallagin, and the glycosides vitexin and rutin (Yun-Lian et al., 2000; Fan et al., 2004; Chyau et al., 2006; Duke, 2008). Additionally, the leaves are rich in flavonoids, tannins, saponins, phytosterols, alkaloids, steroidal glycosides, and phenolic compounds (Punniya and Vijava, 2014). Citronellyl acetate (64.87%) was the predominant compound in the essential oil of Terminalia bentzoë (Gurib-Fakim and Demarne, 1994), while palmitic acid (35.7%) was the major component in T. chebula (Naik et al., 2010). In Terminalia ivorensis,  $\delta$ -3-carene (29.4%) and  $\alpha$ pinene (20.9%) were the most abundant constituents (Ogunwande et al., 2019). Additionally, the essential oils extracted from various parts of T. catappa, including its fruits, leaves, and nuts, have been found to contain several major phytochemicals, such as  $\alpha$ -farnesene (21.3%), hexahydrofarnesyl acetone (12.34%), octadecane (11.7%), hexadecanoic acid (9.5%), dibutyl phthalate (9.1%), 1,2,3-trimethoxy-5-(2propenyl)-benzene (6.6%), neoisothujol (5.8%), 1,2,4-trimethoxy-5-(1-propenyl)-benzene (4.5%), 6,10,14-trimethyl-2-pentadecanoic acid, 1-(2,3,6trimethylphenyl)-(E)-3-buten-2-one, geranyl acetone, hexadecanoic acid (21.0%), 2-ethyl-3,6dimethylpyrazine (19.2%), (Z)-phytol (41.2%), palmitic acid (11.0%), and (E)-nerolidol (4.7%) (Moronkola and Ekundayo, 2000; Wang et al., 2000; Lasekan et al., 2012; Owolabi et al., 2013; Ogunmoye et al., 2020b).

Alternanthera pungens Kunth, a member of the Amaranthaceae family, is commonly known as khaki weed. Locally referred to as "dágunró" in Yoruba-speaking regions of southwestern Nigeria. it is widely distributed across tropical and subtropical areas worldwide (Burkill, 1985). This herbaceous perennial plant typically has prostrate stems, occasionally upright, ranging from 10 to 50 cm in length (Naidu, 2012; Hossain et al., 2018). Livestock generally avoid consuming khaki weed, as it is suspected to be toxic to sheep and pigs, and has been associated with digestive issues and skin problems in cattle (Parsons and Cuthbertson, 2001). Although it is not very palatable to goats, there is no documented evidence of toxicity in them (Simmonds et al., 2000). The plant has demonstrated various pharmacological properties, including spasmogenic, diuretic, anti-HIV, antioxidant, antimicrobial, antibacterial, analgesic, and antidiabetic effects (Petrus and Seetharaman, 2005; Ogundare and Oladejo, 2014). It is also traditionally used to treat gastrointestinal disorders (Adela et al., 2008) and is employed in Mexican folk medicine for managing diarrhea and dysentery (Osuna et al., 2005).

Phytochemical analyses of the plant extracts have revealed the presence of several compounds, including saponins, steroids, alkaloids, triterpenoids,  $\beta$ -spinasterol, glycosides, flavonoids, tannins, phenols, carbohydrates,  $\beta$ carotene, choline, violaxanthin, zeaxanthin, and lutein (Hossain et al., 2018; Gupta and Saxena, 1987; Mourya et al., 2018).

Gas Chromatography-Mass Spectrometry (GC-MS) analysis of the essential oil extracted from A. pungens leaves revealed that it is primarily composed of  $\beta$ -ionone (42.18%) and hexahydrofarnesyl acetone (15.53%), with other constituents present in trace amounts (Ogunmoye et al., 2020a). Similarly, the essential oil from the plant's flowers has been found to contain a variety of compounds, including  $\alpha$ -pinene (7.40%),  $\beta$ -pinene (6.42%), camphene (4.21%), myrcene (3.61%), p-cymene (4.29%), limonene (3.52%),  $\beta$ -ocimene (2.35%), 1,8-cineole (6.28%),  $\alpha$ -thujone (3.62%),  $\alpha$ -borneol (4.46%),  $\alpha$ -curcumene

(2.36%), camphor (5.52%), linalool (6.29%), geraniol (7.42%),  $\alpha$ -terpineol (3.82%), elemol acetate (6.14%), eudesmol (5.38%), azulene (3.16%), along with three unidentified compounds (Gupta and Saxena, 1987).

As a continuation of our investigation into the essential oils of these plants, the present study focuses on evaluating the antioxidant activity of the essential oils extracted from the leaves of Terminalia catappa and Alternanthera pungens using the DPPH assay.

#### **MATERIALS AND METHODS**

#### **Plant materials**

Fresh leaves of Terminalia catappa and Alternanthera pungens were collected from trees located within the premises of Olabisi Onabanjo University, Ago Iwoye, Ogun State, Nigeria. The plant samples were identified and authenticated by Mr. A. O. Adeyemo at the Forestry Research Institute of Nigeria (FRIN), Ibadan, and assigned herbarium numbers FHI-110462 and FHI-110461. Voucher specimens were subsequently deposited at the institute's herbarium. The collected leaves were air-dried at ambient room temperature and then ground into a fine powder for further analysis.

### **Extraction of essential oils**

A total of 200 g of air-dried, pulverized T. catappa leaves and 500 g of A. pungens leaves were subjected to hydrodistillation using a Clevenger-type all-glass apparatus for three hours, following standard protocols (British Pharmacopoeia, 1980; Ogundajo et al., 2016). The essential oils obtained were collected in hexane, transferred into sample bottles, and stored in a refrigerator for subsequent analysis.

Estimation of DPPH Radical Scavenging Activity of Terminalia catappa and Alternanthera pungens.

Stock solutions of each plant extract were prepared in methanol at a concentration of 10 mg/mL. These were further diluted to obtain concentrations of 100, 50, 25, and 12.5 mg/mL. Ascorbic acid served as the standard, prepared in concentrations ranging from 1 to 100 mg/mL. Each dilution was combined with 1 mL of DPPH solution, and the mixtures were kept in the dark at room temperature for 30 minutes. The absorbance was then measured at 517 nm using a T90+ UV/VIS spectrophotometer, following the method outlined by Saleh et al., 2010, with slight modifications. All experiments were conducted in triplicate.

The percentage inhibition was calculated using the following formula:

$$AA = \frac{Ac - As}{Ac} x 100$$

Where

 $A_c =$  the absorbance of control.

 $A_s =$  the absorbance of the test (sample).

AA=the antioxidant activity.

The standard ascorbic acid (vitamin C) was also examined for its antioxidant activity using the same methodology for comparison.

#### RESULTS

The essential oils extracted from the air-dried leaves varied in appearance, ranging from a cloudy light yellow to a cloudy white liquid, each possessing a strong aroma. The extraction yielded 0.25% for *Terminalia catappa* and 0.40% for *Alternanthera pungens*. Tables 1.0 and 2.0 present the percentage inhibition values of the essential oils from *T. catappa* and *A. pungens*, respectively, compared to a standard reference. Additionally, Figures 1.0 and 2.0 illustrate the percentage inhibition graphs representing the antioxidant activity of the essential oils from *T. catappa* and *A. pungens*, respectively.

#### **DISCUSSION AND CONCLUSION**

Based on the data presented in Table 1.0 and Figure 1.0, the essential oil demonstrated varying levels of percentage inhibition across different concentrations ranging from 100 mg/mL to 12.5 mg/mL. Specifically, the inhibition percentages were 54.22% at 100 mg/mL, 82.62% at 50 mg/mL, 84.67% at 25 mg/mL, and 79.07% at 12.5 mg/mL. These results indicate that the antioxidant activity fluctuated with concentration, showing a moderate level of efficacy with inhibition values between 54.22% and 84.67%. The lowest inhibition was recorded at 100 mg/mL (54.22%), while the highest occurred at 25 mg/mL (84.67%). Overall, the essential oil displayed significant DPPH free radical scavenging activity that was generally concentration-dependent, with the exception of the lowest concentration (12.5 mg/mL), which still maintained a high inhibition rate of 79.07%. These findings confirm the antioxidant potential of the oil

According to the data in Table 2.0 and Figure 2.0, the essential oil exhibited varying levels of percentage inhibition at concentrations ranging from 100 mg/mL to 12.5 mg/mL. The observed inhibition values were 73.87% at 100 mg/mL, 43.90% at 50 mg/mL, 68.13% at 25 mg/mL, and 73.46% at 12.5 mg/mL. These results indicate noticeable variation in antioxidant activity across the tested concentrations. The oil demonstrated moderate antioxidant potential, with inhibition percentages ranging from 43.90% to 73.87%. The lowest activity was recorded at 50 mg/mL (43.90%), while the highest was at 100 mg/mL (73.87%). Unlike the standard, the free radical scavenging activity of this essential oil did not follow a consistent concentration-dependent pattern, as shown by the irregular inhibition values. Nonetheless, the results confirm the presence of antioxidant activity based on the DPPH assay.

The essential oil extracted from Terminalia catappa exhibited strong antioxidant activity, with inhibition values ranging from 54.22% to 84.67%, the highest being 84.67% at a concentration of 25 mg/mL. In comparison, Alternanthera pungens showed moderate antioxidant activity, achieving a maximum inhibition of 73.87% at 100 mg/mL. In this study, the observed reduction in DPPH color from purple to yellow across various concentrations indicates the presence of electrondonating compounds within the oils, which are capable of neutralizing free radicals by converting them into more stable molecules. These findings confirm the antioxidant potential of the leaf essential oils, aligning with earlier studies on T. catappa leaf extracts (Punniva and Vijaya, 2014; Rajesh et al., 2015).

In conclusion, the essential oils demonstrated moderate to strong antioxidant activity in comparison to the radical scavenging potential of the ascorbic acid standard. Notably, the oils extracted from Terminalia catappa and Alternanthera pungens appear to be promising and rich sources of natural antioxidants. This indicates their potential use as antioxidant agents in industries such as food, pharmaceuticals, and cosmetics. However, further antioxidant assays are recommended to confirm and support the results obtained through the DPPH method.

The authors express their sincere gratitude to Miss Hammed Rashidat Oluwakemi and Miss Oyebade Doyinsola Christianah for their valuable assistance during the extraction of the oil samples.

Table 1.0: % Inhibition of *T. catappa* leaves essential oil and known antioxidant on DDPH

Concentration (mg/mL)	Oil sample	Ascorbic acid	
100 mL	54.22%	91.75%	
50 mL	82.62%	89.19%	
25 mL	84.67%	87.27%	
12.5 mL	79.07%	86.18%	



Fig. 1.0: Graph showing the antioxidant activity of the essential oil of T. catappa

Table	2.0:	%	Inhibition	of <i>A</i> .	pungens	leaves
essential oil and known antioxidant on DDPH						

Concentration (mg/mL)	Oil sample	Ascorbic acid	
100 mL	73.87%	91.5%	
50 mL	43.90%	89.19%	
25 mL	68.13%	87.27%	
12.5 mL	73.46%	74.41%	



Fig. 2.0: Graph showing the antioxidant activity of the essential oil of A. pungens

#### Ogunmoye A. O. et al.: DPPH activities of T. catappa and A. pungens air-dried leaves

#### REFERENCES

- Adela A, Hortencia D, Leyvert D, Guadalupe H and Andrés N. (2008). Investigation of *Alternanthera repens* and *Bidens odorata* on gastrointestinal disease. Fitoterapia 07:001-007 Adv. Exp. Med. Biol. Advances in Experimental Medicine and Biology 595: 1–75. doi: 10.1007/978-0-387-46401-5\_1. ISBN 978-0-387-46400-8. PMID 17569205.
- Agu C.M and Menkiti M.C. (2017). Effects of natural antioxidants on the essential properties of modified *Terminalia catappa* L. kernel oil: A possible substitute for mineral oil transformer fluid. Biofuels.
- Aruoma O.I. (1998). Free radicals, oxidative stress, and antioxidants in human health and disease. *J. Am. Oil Chem. Soc.* **75:**199-212.
- British Pharmacopoeia 1980. Her Majesty;s Stationary Office, Atlantic House, Holborn Viaduct, London, England Vol. II, pp. 109.
- Burkill H.M. (1985). The useful plants of west tropical Africa. *Royal Botanic Gardens*, Kew, UK. **3**.
- Chu S.C, Yang S.E, Liu S.J, Kuo W.H and Chang Y.Z. (2007). *Terminalia catappa* L. leaves on lung cancer Cells. *Food and Chemical Toxicology*. 45(7):1194-1201.
- Chyau C.C, Ko P.T and Mau J.L. (2006). Antioxidant properties of aqueous extract from *Terminalia catappa* leaves. *Food Sci. Technol.* (LWT). 39:1099-1108
- Duke JA. Phytochemical and Ethnobotanical Databases. 2008. (online database).
- Elizabeth K.M. (2005). Antimicrobial activity of *Terminalia* bellerica.Indian J. Clin. Biochem. **20(2)**: 150-153.
- Ezeokonkwo C.A and Dodson W.L. (2004). The potential of *Terminalia catappa* (tropical almond) seed as a source of dietary protein. *J. Food Qual.* **27(3)**:207–219.
- Fan Y.M, Xu L.Z, Gao J, Wang Y, Tang X.H, Zhao X.N and Zhang Z.X. (2004). Phytochemical and antiinflammatory studies on *Terminalia catappa*. *Fitoterapia*. 75:253–260.
- Gbile, Z.O. (1984). Vernacular Names of Nigerian Plants (Yoruba), Forestry Research Institute of Nigeria.
- Gupta R.K and Saxena V.K. (1987). Volatile constituents from the flowers of *Alternanthera pungens* HBK (Amaranthaceae). Indian Perfume.31(4):366-369.
- Gurib-Fakim A and Demarne F. (1994). Essential oil of *Terminalia bentzoë* (L.) L. f. subsp. Rodriguesensis Wickens. *Journal of Essential Oil Research*. 6(5):533-534.

- Hossain M.D.S, Rana M.D.S, Rajibul Islam and Mahmudul Islam AFM. (2018). Phenolic content analysis and evaluation of antinociceptive, antioxidant, antiinflammatory potential of *Alternanthera pungens* KUNTH. Journal of Bangladesh Academy of Sciences.42(2):129-136.
- Houghton, P. (1995). The role of plants in traditional medicine and current therapy. *J Alter Comple Med.* 1:131-143.
- Isah J.J. (2021). Drying effect on the isolation, characterization and antioxidant potentials of leaf essential oils of *Ocimum gratissimum* harvested at 10:00am in a day.*Journal of Experimental Research*. **9(3)**:39-48
- Kamatou G.PP. and Viljoen A.M. (2010). A review of the application and pharmacological properties of αbisabolol and α-bisabolol-rich oils. J. Am. Oil Chem. Soc. 87, 1-7.
- Ko T.F, Weng Y.M and Chiou R.Y.Y. (2002). Squalene content and antioxidant activity of *Terminalia catappa* leaves and seeds. J. Agric. Food Chem. 50:5343-5348
- Kumar R.S, Sivakumar T, Sunderam R.S, Gupta M, Mazumdar U.K, Gomathi P, Rajeshwar Y, Saravanan S, Kumar M.S, Murugesh K and Kumar K.A. (2005). Antioxidant and antimicrobial activities of *Bauhinia* racemosa L. stem bark. *Brazil J Med Biol Res*:1015–1024.
- Lasekan O, Alfi K and Abbas K.A. (2012). Volatile compounds of roasted and steamed Malaysian tropical almond nut (*Terminalia catappa* L.). *Int. J. Food Prop.* 15(5):1120-1132.
- Lin C.C and Kan W.S. (1990). Medicinal plants used for the treatment of hepatitis in Taiwan. *Am. J. Chinese Med.* 18:35-43.
- Maestri D.M, Nepote V, Lamarque A.L and Zygadlo J.A. (2006). Natural products as antioxidants. In Phytochemistry: Advances in Research; Imperato, F., Ed.; Research Signopost: Kerala, India. pp. 105-135.
- Mety S.S and Mathad P. (2011). Antioxidative and free radical scavenging activities of Terminalia species. *Int. Res. J. Biotech.* **2(5)**:119-127.
- Miguel M.G. (2010). Antioxidant activity of medicinal and aromatic plants. Flavour Fragr. J. 25, 291-312.
- Moronkola D.O and Ekundayo O. (2000). Chemical constituents in the fruit essential oil of *Terminalia catappa* Linn (almond fruits). *J. Trop. For. Res.* **16(1)**:72-79.
- Mourya P, Singh G, Jain N and Gupta M.K. (2018). *Invitrostudies on inhibition of alpha amylase and alpha* glucosidase by plant extracts of *Alternanthera pungens KUNTH. Journal of Drug Delivery & Therapeutics.* 8(6-A):64-68.

- Naidu V.S.G.R. (2012). Hand Book on Weed Identification. Directorate of Weed Science Research, Jabalpur, India. pp. 354
- Naik D.G, Puntambekar H and Anantpure P. (2010). Essential oil of *Terminalia chebula* fruits as a repellent for the Indian honeybee Apisflorea. *Chem. Biodiver.* 7(5):1303-1310.
- Nair R and Chanda S. (2008). Antimicrobial activity of *Terminalia catappa*, *Manilkara zapota* and *Piper betel* leaf extract. *India J. Pharm. Sci.* 70(30):390-393
- Ogundajo A.L, Nnaemeka C.O, Olawunmi R.O and Ogunwande I.A. (2016). Chemical Constituents of Essential oil of Ethretia cymosa Thonn. *British Journal* of Applied Science and Technology. **14(4)**:1-6.
- Ogundare A.O and Oladejo B.O. (2014). Antibacterial activities of the plant extract of Alternanthera pungens. *European Journal of Botany Plant Sciences and Phytology*. **1(4)**:1-7.
- Ogunmoye A.O, Atewolara-Odule O.C, Olubomehin O.O, Ogundare S.A, and Yussuf S.T. (2020a). The chemical constituents' of the leaf essential oil of *Alternanthera pungens* (KUNTH). *African Journal of Science and Nature*. 10:131-136.
- Ogunmoye A.O, Olubomehin O.O, Atewolara-Odule O.C, Ogundare S.A, and Yussuf S.T. (2020b). GC-MS Analysis of the volatile constituents from the air-dried leaves of *Terminalia catappa* (LINNAEUS).*FUW Trends in Science & Technology Journal.* **5(3)**: 948 – 951.
- Ogunwande I.A, Ascrizzi R, and Guido F. (2019). Essential oil composition of *Terminalia ivorensis* A. Chev. flowers from northern Nigeria. *Trends in Phytochem. Res.* **3(1)**:77-82
- Orwa C, Mutua A, Kindt R, Jamnadass R, and Anthony S. (2009). Agroforestree Database: A Tree Reference and Selection Guide. Version 4.0.
- Osawa T, Kavakishi S, Namiki M, Kuroda Y, Shankai O.M, and Waters M.D. (1990). Antimutagenesis and Antimutagenesis mechanism, Edn 11, New York plenum. 139-153.
- Osuna L, Tapia-Pérez M, Jiménez-Ferrer J, Carrillo-Quiróz B, and Silva-Sánchez J. (2005). Screening of *Alternanthera repens., Boerhavia coccinea., Flaveria trinervia.*, Tournefortia.
- Owolabi M.S, Lawal O.A, Ogunwande I.A, Hauser R.M, William N, and Setzer W.N. (2013). Chemical composition of the leaf essential oil of *Terminalia catappa* L. growing in southwestern Nigeria. *Am. J. Essential Oils and Nat. Prod.* **1(1)**:51-54.

- Parsons W.T, and Cuthbertson E.G. (2001). *Noxious Weeds of Australia*. Second Edition ed. Collingwood, Victoria: CSIRO Publishing. pp. 705.
- Petrus AJA and Seetharaman TR. (2005). Antioxidant flavone C-biosides from the aerial parts of *Alternanthera pungens. Indian Journal of Pharmaceutical Sciences.* **67(2)**:187-193.
- Pham-Huy L.A, He H and Pham-Huyc C. (2008). Free Radicals, Antioxidants in Disease and Health. *International Journal of Biomedical Science*. **4(2)**:89-96.
- Punniya P.K and Vijaya A.A.(2014). Phytochemical analysis and *in vitro* antioxidant activity of *Terminalia catappa*. *World Journal of Pharmaceutical Sciences*. 2(11):1495-1498.
- Rajarajan S, Asthana M and Shanthi, G. (2010). The in vitro bactericidal activity of lyophilized ethanolic extract of Indian almond (*Terminalia catappa* Linn) fruit pulp on two pathogenic bacteria from subgingival plaques. *Indian J. Nat. Prod. and Resou.* 1(4):466-469.
- Rajesh B.R, Potty V.P, Prabha K.C, Miranda M.T.P and Sreelekshmy S.G. (2015). Antioxidant and antimicrobial activity of leaves of *Terminalia catappa* and *Anacardium occidentale*: A comparative study. *Journal of Pharmacognosy and Phytochemistry*. **4(1)**: 79-82
- Rao NK and Nammi S. (2006). Antidiabetic and renoprotective effects of the chloroform extract of *Terminalia chebularetz*, Seeds in streptozotoc induced diabetic rats. BMC: CAM. 6:17-17.
- Saleh M.A, Clark S, Woodward B, and Deolu-Sobogun S.A. (2010). Antioxidant and Free Radical Scavenging Activities of Essential oils. *Ethnicity and Disease*. 20:78-82.
- Simmonds H, Holst P, and Bourke C. (2000). *The Palatability, and Potential Toxicity of Australian Weeds to Goats.* Canberra: Rural Industries Research and Development.
- Untwal L.S and Kondawar M.S. (2006). Use of *Terminalia catappa* fruit extract as an indicator in acid-base titration. *Indian J. Pharmac. Sci.* 68(3):399-401.
- Wang .HF, Ko P, Chyau C.C, Mau J.L and Kao M.D. (2000). Composition and antioxidative activity of essential oils from *Terminalia catappa* L. leaves. *Taiwanes eJournal* of Agric. Chem. and Food Sci. 38(1):27-35.
- Willcox J.K, Ash S.L and Catignani G.L. (2004). Antioxidants and prevention of chronic disease. Critical Reviews in Food Science and Nutrition. 44:275-295.
- Yun-Lian L, Yueh-Hsiung K, Ming-Shi S, Chien-Chih C and Jun-Chih O. (2000). Flavonoid glycosides from *Terminalia catappa L.J. Chin. Chem. Soc.* 47:253-256.

249