**Original Research Article**

# Proximate analysis and phytochemical profile of *Brachystegia eurycoma* leaves


## ABSTRACT

**Aim:** This study investigated the proximate and phytochemical composition of *Brachystegia eurycoma* leaves.

**Methods:** Crude ethanol extract of *B. eurycoma* leaves was obtained by cold extraction method. AOAC method was used for proximate analysis. Phytochemical profiling was done with qualitative phytochemical evaluation and gas chromatography-mass spectrometry (GC/MS) analytical method. Matching and interpretation of the spectral was done with the National Institute standard and Technology (NIST05) library.

**Result:** The proximate analysis result showed *B. eurycoma* leaves to be abundant in parameters evaluated in the order of 31.47±0.43% Carbohydrate > 15.15±0.04% Ash > 14.45±0.15 crude fibre > 13.83±0.32 protein > 13.14±0.22 moisture > 1.97±0.01 fat. Qualitative phytochemical analysis detected alkaloid, saponin, tanin, diterpenes, phenol, quinine, flavonoid, protein, xanthoprotein and cardiac glycoside in the leaves of *B. eurycoma*. GC/MS data showed that the prevailing bioactive compounds in ethanol leaf extract of *B. eurycoma* were 3-O-Methyl-d-glucose (13.23%), cis-9-Hexadecenal (10.40%), Desulphosinigrin (10.34%), Phytol (7.58%), Hydroquinone (7.23%), n-Hexadecanoic acid (6.61%), Oleoyl chloride (6.10%), 9,12-Octadecadienoic acid (Z,Z)- (5.89%), Hexadecanoic acid, (2.97%), Benzofuran, 2,3-dihydro- (1.94%), Hexadecanoic acid, 2-hydroxy-1-(hydroxymethyl) (1.92%).

**Conclusion:** This result reveals the potentials of *B. eurycoma* leaves in food, pharmaceutical, cosmetic and nutraceutical industry.

**Keywords:** *Brachystegia eurycoma*, Gas chromatography mass spectrometry (GC-MS), phytochemical, nutraceutical, Desulphosinigrin, drug discovery.

## INTRODUCTION

Plants contain a wide range of bioactive chemical substances (flavonoids, alkaloids, steroids, trepenoids, phenolic acids, tannins, saponins among others) that exhibits therapeutic, physiological and biochemical effects in human body [1-4]. Scientific research resulting in phytocomponent profiling, isolation, purification and characterization of phytochemicals has led to the discovery of drug candidates, production of active drugs, supplements and food additives used in the treatment and management of different ailments all over the world [5-7].

Proximate analysis is an important procedure to determine the overall composition, nutritional status, quality and energy value of any ingredient intended for use as food [8]. Preliminary phytochemical screening is used to identify the classes of bioactive constituents present in plant [9]. Gas Chromatography Mass Spectroscopy is a technique which is used to separate and identify drug based on retention time and fragmentation pattern. The fragmentation pattern for a drug is unique and is therefore an identifying characteristic of the drug [10, 11]. GC-MS studies have been increasingly applied for the analysis of medicinal plants in recent year as this technique has proved to be a valuable method for the analysis of non-polar components and volatile essential oil, fatty acid, lipids and alkaloids [12].

*Brachystegia eurycoma* is a dicotyledonous leguminous tress belonging to the family *Fabaceae*. It is found in the swamps and rainforest of south, east and western Nigeria. It is called “Achi” by

Igbos, “Ekalado” by the yorubas, “Taura” by the hausas, “Okweri” by the Edos and “Apaupan” by the Ijaw. In the eastern part of Nigeria, its seeds are used as thickener in preparation of local soups particularly egusi and ogbono soup, while its wound healing and inflammatory properties have been mentioned in literature [13].

The leaf, bark and root of the plant are used in ethno medicines for the treatment of various diseases including malaria, diabetes, rheumatism, hypertension, kidney problem, asthma, tuberculosis, bronchitis, catarrh and sore throat [14-16].

Research has shown the seed, stem bark and stem gum as well as phytocomponents isolated from them to be antiinflamatory, antibacterial, analgesic, antioxidant, antimicrobial, anti cancer and anti diarrhea [17-21]. Several researches have been done on the seeds, stem bark and stem gum of *B. eurycoma* with a little attention on the leaves which despite its medicinal use with other parts of the plant is not eaten as food. Hence this research to evaluate the nutrition and phytochemical components of *B. eurycoma* leaves with the aim of unveiling its possible nutritional and nutraceutical potentials.

## MATERIALS AND METHODS

### Plant material

*B. eurycoma* leaves were collected from Abakiliki, Ebonyi State, Nigeria, authenticated at the University of Port Harcourt Herbarium, Port Harcourt, Nigeria. The leaves were rinsed in distilled water, air-dried and ground into powder.

**Preparation of ethanol extract**

The leaf powder was soaked in ethanol (70%) for 48 hours with occasional shaking. It was filtered and the filtrate concentrated after eliminating the ethanol using rotary evaporator.

**Determination of proximate composition**

The amount (%) of moisture, ash, lipid, protein, fiber, carbohydrates as well as energy level in air dried *B. eurycoma leaf* was determined in triplicate using the method of AOAC [22].

**Preliminary phytochemical determination**

The concentrated leaves extract of *B. eurycoma* were analyzed for the presence of saponins, flavonoids, tanins, alkaloids, phenols, quinines, protein, xanthoprotein, cardiac glucoside, Coumarin, steroids, diterpene and Anthraquinone using standard methods as described by Fafowora [23], Evans [24] and Harborne [25].

**Gas Chromatograph-Mass Spectroscopy (GC-MS) analysis**

GC-MS was performed on a system consisting of GC2010 gas chromatograph and a Shimadzu QP2010 ultra quadrupole mass spectrometer equipped with a DB-Wax fused silica capillary column (30m x 0.25mm ID x 0.25µm df, composed of silphenylene polymer). Initial temperature was programmed at 50$^{o}$C, held for two minutes. It was increased to 300$^{o}$C with the rate of 6.5$^{o}$C/min and held for ten minutes. The temperature of the injector and detector were set up to 280$^{o}$C and 300$^{o}$C, respectively. Helium gas was used as a carrier gas. 1 µl of the fractions was diluted in 200 µl dichloromethane and then injected into the GC-MS [26, 27].

**Identification of spectral**

Interpretation of mass-spectrum was done by comparing the mass spectrum of the unknown component with spectrum of known components in the National Institute Standard and Technology (NIST) database to ascertain the name, molecular weight and structure of the components of the test materials.

**RESULT**

The result of proximate analysis shows that the leaves of *B. eurycoma* to contain 31.47±0.43% carbohydrates, 15.15±0.04% ash, 14.45±0.15% crude fiber, 13.83±0.32% protein, 13.14±0.22% moisture, 1.97±0.01% fat and 198.87±0.56 Kcal/100g (Table 1).

**Table 1.** Proximate composition of *B. eurycoma* leaves.

| **Parameter** | **Unit** | ***B. eurycoma*** |
|---|---|---|
| Carbohydrate (NFE) | % | 31.47±0.43 |
| Ash | % | 15.15±0.04 |
| Crude fibre | % | 14.45±0.15 |
| Protein | % | 13.83±0.32 |
| Moisture | % | 13.14±0.22 |
| Fat | % | 1.97±0.01 |
| Energy level | Kcal/100g | 198.87±0.56 |

Qualitative phytochemical analysis revealed the presence of alkaloid, saponin, tanin, diterpenes, phenol, quinine, flavonoid, protein, xanthoprotein and cardiac glycoside in the leaves of *B. eurycoma.* Coumarin, steroids and anthraquinone were not detected (Table 2).

**Table 2.** Qualitative phytochemical evaluation of *B. eurycoma* leaves.

| **Phytochemical components** | **Type of test** | **Ethanol leaf extract of *B. eurycoma*** |
|---|---|---|

| | | | |
|---|---|---|---|
| 1 | Alkaloids | Mayer's Test | + |
| | | Wanger's test | + |
| | | Dragendroff's test | + |
| 2 | Phenol | Ferric Chloride | + |
| 3 | saponin | Frothing | + |
| 4 | Tannin | Ferric Chloride | + |
| 5 | Coumarin | alcoholic sodium hydroxide | - |
| 6 | Quinine | potassium hydroxide | + |
| 7 | Anthraquinone | Borntrager's | - |
| 8 | steroids | Libermann- Burchard | - |
| 9 | Protein | Million's | + |
| 10 | Xanthoprotein | General test | + |
| 11 | Cardiac glycoside | Keller kiliani | + |
| 12 | Flavonoid | sodium hydroxide | + |
| 13 | Diterpene | Copper acetate | + |

The GC-MS chromatogram obtained from ethanol extract of *B. eurycoma* shows the presence of 24 distinct peaks (figure 1).

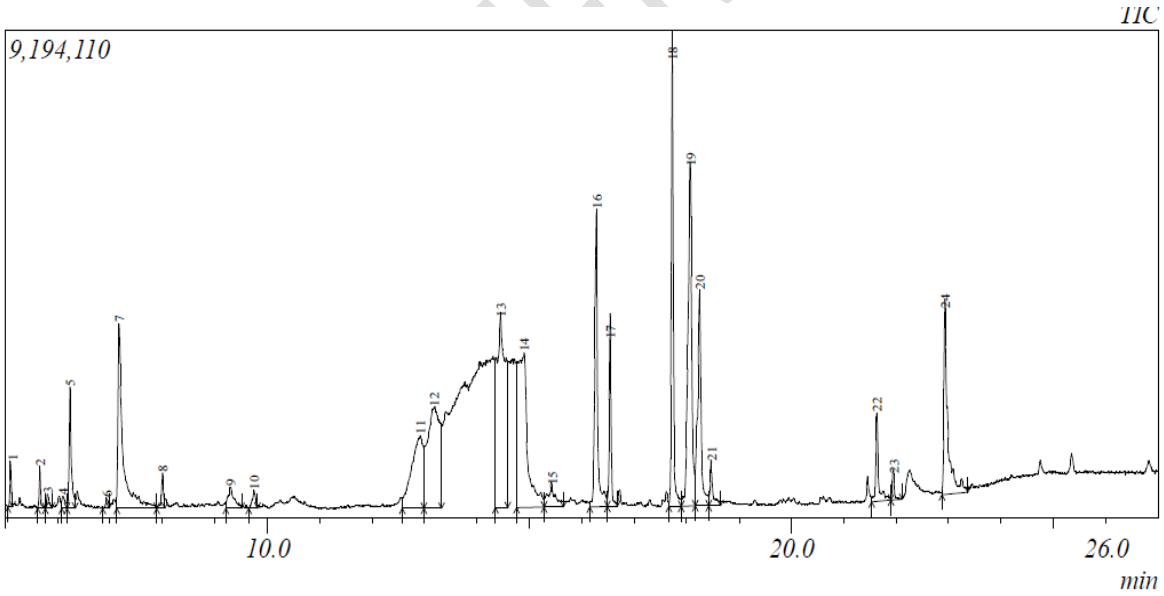



**Figure 1. GC-MS chromatogram of ethanol leaf extract of *B. eurycoma***

The 24 phytochemicals identified in the order of their peaks are presented in figure 2.

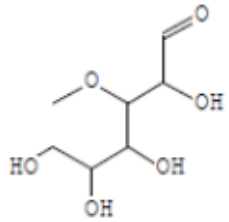
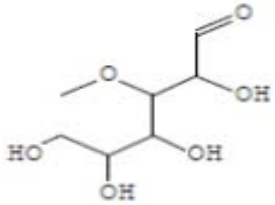
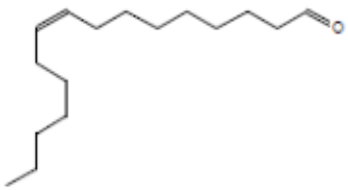
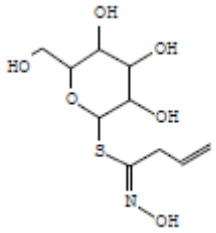
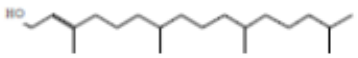
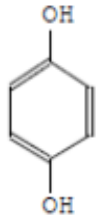
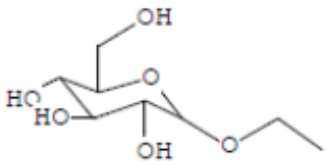
Peak Report TIC

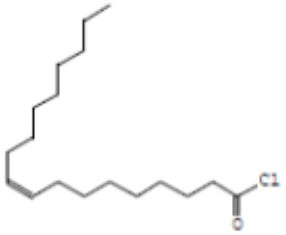
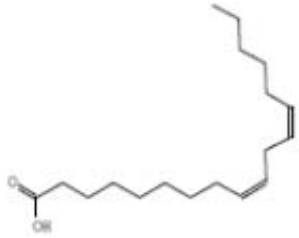
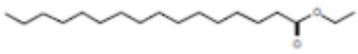
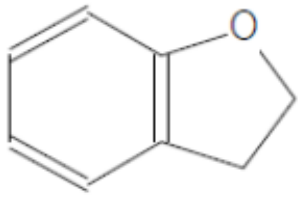
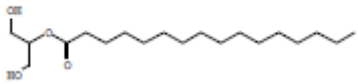
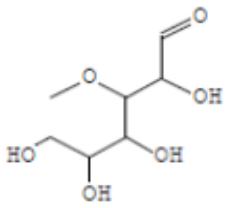
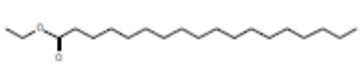
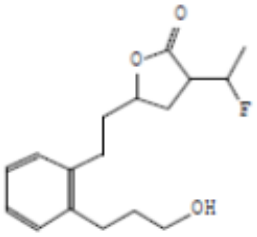
| Peak# | R.Time | I.Time | F.Time | Area | Area% | Height | Height% | A/H | Mark | Name |
|---|---|---|---|---|---|---|---|---|---|---|
| 1 | 5.102 | 5.050 | 5.142 | 1572759 | 0.50 | 868088 | 1.60 | 1.81 | | Furan-3-carboxaldehyde, 2-methoxy-2,3-dil |
| 2 | 5.660 | 5.625 | 5.767 | 1578299 | 0.50 | 785361 | 1.45 | 1.99 | | 4H-Pyran-4-one, 2,3-dihydro-3,5-dihydroxy |
| 3 | 5.806 | 5.767 | 5.883 | 821421 | 0.26 | 271589 | 0.50 | 3.02 | V | 2-Cyclopenten-1-one, 3-ethyl-2-hydroxy- |
| 4 | 6.100 | 6.075 | 6.183 | 986554 | 0.31 | 233514 | 0.43 | 4.22 | V | Hydroquinone |
| 5 | 6.237 | 6.183 | 6.333 | 6085352 | 1.94 | 2283993 | 4.22 | 2.66 | V | Benzofuran, 2,3-dihydro- |
| 6 | 6.935 | 6.867 | 6.983 | 618211 | 0.20 | 190340 | 0.35 | 3.25 | V | Pipridine, 1-chloroacetyl- |
| 7 | 7.170 | 7.125 | 7.892 | 22647479 | 7.23 | 3485442 | 6.44 | 6.26 | V | Hydroquinone |
| 8 | 8.002 | 7.892 | 8.042 | 1741729 | 0.56 | 675050 | 1.25 | 2.58 | V | Benzene, 2-methoxy-1,3,5-trimethyl- |
| 9 | 9.299 | 9.208 | 9.508 | 3093625 | 0.99 | 401110 | 0.74 | 7.71 | V | Tetrahydrofuran-2-one, 3-[1-fluoroethyl]-5- |
| 10 | 9.747 | 9.650 | 9.817 | 1409666 | 0.45 | 338811 | 0.63 | 4.16 | | 9-[2-Deoxy-.beta.-d-ribohexopyranosyl]pur |
| 11 | 12.916 | 12.575 | 12.983 | 20727848 | 6.61 | 1374145 | 2.54 | 15.08 | V | Ethyl .alpha.-d-glucopyranoside |
| 12 | 13.177 | 12.983 | 13.325 | 32406722 | 10.34 | 1895570 | 3.50 | 17.10 | V | Desulphosinigrin |
| 13 | 14.454 | 14.350 | 14.583 | 41452388 | 13.23 | 3597376 | 6.65 | 11.52 | V | 3-O-Methyl-d-glucose |
| 14 | 14.900 | 14.750 | 15.283 | 38929701 | 12.42 | 2879052 | 5.32 | 13.52 | V | 3-O-Methyl-d-glucose |
| 15 | 15.433 | 15.283 | 15.650 | 4468492 | 1.43 | 429751 | 0.79 | 10.40 | V | 3-O-Methyl-d-glucose |
| 16 | 16.281 | 16.150 | 16.483 | 20727534 | 6.61 | 5542816 | 10.24 | 3.74 | V | n-Hexadecanoic acid |
| 17 | 16.547 | 16.483 | 16.683 | 9309686 | 2.97 | 3395336 | 6.27 | 2.74 | V | Hexadecanoic acid, ethyl ester |
| 18 | 17.732 | 17.667 | 17.900 | 23743294 | 7.58 | 8683680 | 16.05 | 2.73 | V | Phytol |
| 19 | 18.072 | 17.900 | 18.167 | 32593772 | 10.40 | 6241435 | 11.53 | 5.22 | V | cis-9-Hexadecenal |
| 20 | 18.250 | 18.167 | 18.417 | 18472090 | 5.89 | 4009206 | 7.41 | 4.61 | V | 9,12-Octadecadienoic acid (Z,Z)- |
| 21 | 18.473 | 18.417 | 18.650 | 3301278 | 1.05 | 767072 | 1.42 | 4.30 | V | Octadecanoic acid, ethyl ester |
| 22 | 21.634 | 21.533 | 21.900 | 6010628 | 1.92 | 1633519 | 3.02 | 3.68 | V | Hexadecanoic acid, 2-hydroxy-1-(hydroxyn |
| 23 | 21.950 | 21.900 | 22.117 | 1542228 | 0.49 | 538818 | 1.00 | 2.86 | V | Bis(2-ethylhexyl) phthalate |
| 24 | 22.933 | 22.867 | 23.367 | 19114576 | 6.10 | 3592028 | 6.64 | 5.23 | V | Oleoyl chloride |
| | | | | 313355332 | 100.00 | 54113102 | 100.00 | | | |

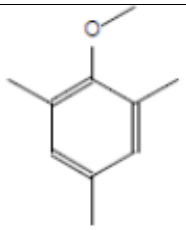
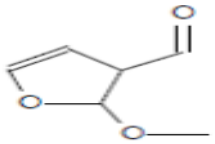
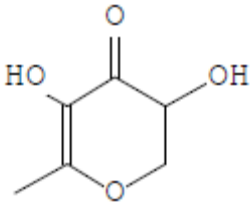
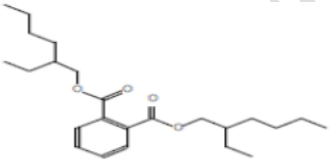
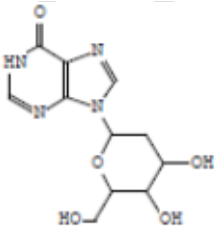
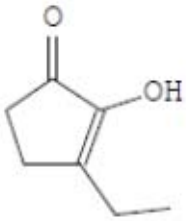
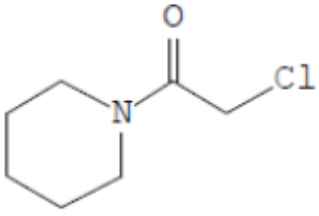
**Figure 2.** Phytochemicals present in ethanol leaf extract of *B. eurycoma*

The first compound identified with less retention time (5.102 min) was Furan-3-carboxaldehyde 2-methoxy-2,3-dihydro- whereas Oleoyl chloride was the last compound which took longest retention time (22.933 min) to identify. The phytochemicals identified in order of most abundant to least abundant based on percentage peak are presented in table 3, showing their molecular formular and structure.

**Table 3.** Molecular formula and structure of phytochemicals identified in *B. eurycoma*.

| | Name of compound | Molecular formula | Structure | Molecular weight | Peak Area (%) |
|---|---|---|---|---|---|
| 1 | 3-O-Methyl-d-glucose | $C_7H_{14}O_6$ |  | 194 | 13.23 |
| 2 | 3-O-Methyl-d-glucose | $C_7H_{14}O_6$ |  | 194 | 12.42 |
| 3 | cis-9-Hexadecenal ; 9-Hexadecenal, (Z)- | $C_{16}H_{30}O$ |  | 238 | 10.40 |
| 4 | Desulphosinigrin | $C_{10}H_{17}NO_6S$ |  | 279 | 10.34 |
| 5 | Phytol | $C_{20}H_{40}O$ |  | 296 | 7.58 |
| 6 | Hydroquinone | $C_6H_6O_2$ |  | 110 | 7.23 |
| 7 | n-Hexadecanoic acid | $C_{16}H_{32}O_2$ |  | 256 | 6.61 |
| 8 | Ethyl .alpha.-d-glucopyranoside | $C_8H_{16}O_6$ |  | 208 | 6.61 |

| | | | | | |
|---|---|---|---|---|---|
| 9 | Oleoyl chloride | $C_{18}H_{33}C_{lO}$ |  | 300 | 6.10 |
| 10 | 9,12-Octadecadienoic acid (Z,Z)- | $C_{18}H_{32}O_2$ |  | 280 | 5.89 |
| 11 | Hexadecanoic acid, ethyl ester ; | $C_{18}H_{36}O_2$ |  | 284 | 2.97 |
| 12 | 2,3-dihydro-Benzofuran Coumaran | $C_8H_8O$ |  | 120 | 1.94 |
| 13 | Hexadecanoic acid, 2-hydroxy-1-(hydroxymethyl)ethyl ester<br><br>Palmitin, 2-mono- | $C_{19}H_{38}O_4$ |  | 330 | 1.92 |
| 14 | 3-O-Methyl-d-glucose | $C_7H_{14}O_6$ |  | 194 | 1.43 |
| 15 | Octadecanoic acid, ethyl ester | $C_{20}H_{40}O_2$ |  | 312 | 1.05 |
| 16 | Tetrahydrofuran-2-one, 3-[1-fluoroethyl]-5-[[2-hydroxypropyl]benzene ethyl- | $C_{17}H_{23}FO_3$ |  | 294 | 0.99 |

| | | | | | |
|---|---|---|---|---|---|
| 17 | Benzene, 2-methoxy-1,3,5-trimethyl- | $C_{10}H_{14}O$ |  | 150 | 0.56 |
| 18 | Furan-3-carboxaldehyde 2-methoxy-2,3-dihydro- | $C_6H_8O_3$ |  | 128 | 0.50 |
| 19 | 4H-Pyran-4-one 2,3-dihydro-3,5-dihydroxy-6-methyl- | $C_6H_8O_4$ |  | 144 | 0.50 |
| 20 | Bis(2-ethylhexyl) phthalate | $C_{24}H_{38}O_4$ |  | 390 | 0.49 |
| 21 | 9-[2-Deoxy-.beta.-d-ribohexopyranosyl]purin-6(1H)-one | $C_{11}H_{14}N_4O_5$ |  | 282 | 0.45 |
| 22 | Hydroquinone | $C_6H_6O_2$ |  | 110 | 0.31 |
| 23 | 2-Cyclopenten-1-one 3-ethyl-2-hydroxy- | $C_7H_{10}O_2$ |  | 126 | 0.26 |
| 24 | Piperidine, 1-chloroacetyl- | $C_7H_{12}C_lNO$ |  | 161 | 0.20 |

## DISCUSSION

The findings from proximate analysis of this study shows *B. eurycoma* leaves to have the abundance of the parameters evaluated to be in Carbohydrate > Ash > crude fibre > protein > moisture > fat order. This shows *B. eurycoma leaves* are a better source of carbohydrates and fiber than proteins. Animals depend on carbohydrates among other macromolecules for generation of energy and some intermediates required for certain biological processes and the sustenance of life. The amount of carbohydrates (31.47%) is appreciable and similar to popular edible vegetables like *Celusia argenta (32.80%)*, *Corchorus olitorius* (31.30%) [29]. The ash content of the leaves (15.15%) is considerable showing that the leaves contain important mineral elements as the ash content of any sample is an index of mineral content.

The RDA of fiber is 19-25% for children, 21-38% for adult, 28% for pregnant mothers and 29% for breast-feeding mothers [30]. With crude fibre of 14.45%, *B. eurycoma* leaves will make a poor source of dietary fiber in human nutrition. Plant foods that provide more than 12% of its calorific value from protein is considered good source of protein [31]. *B. eurycoma* leaves meet this requirement, making it a good source of protein. Moisture content was lower when compared with that of common vegetable such as *Telfairia occidentalis 98%,* Talinum triangulare 91%, moringa oleifera 87% and vernonia amygdalinan 87% [32] but low moisture content is indicative of a longer shelf life. It is a general observation that leafy vegetables have low lipid content [31]. The low lipid content of *B. eurycoma* leaves is in agreement with this observation.

Preliminary phytochemical screening detected the presence of alkaloid, saponin, tanin, diterpenes, phenol, quinine, flavonoid, protein, xanthoprotein and cardiac glycoside. Alkaloids are diuretic in nature, they affect the nervous system and reduce appetite [33]. The ingestion of saponins as a part of the human diet have been linked with a variety of effects on health, including reducing blood cholesterol levels. They have also been reported to have pharmacological activities such as anti-inflammatory, antifungal, antibacterial, anti-parasitic, anti-cancer and antiviral activities [34]. Tannins are antiinflammatory, antidiarrheal, haemostatic, antiviral, antibacterial, antidiarrheal and antihemorrhoidal compounds which has been reported to relief sore throat, fatigue and skin ulcer [35]. Diterpenes are antimicriobial and anti-inflammatory [36]. Quinine has many medicinal applications due to its fever-reducing, painkilling and anti-inflammatory properties [37]. Many flavonoids have been shown to have antioxidative activity, free radical scavenging capacity, antihypertensive, hepatoprotective, anti-inflammatory, antiviral and anticancer activities [38]. Cardiac glycosides have cardiotonic activity, are antiviral, anticancer and antiproliferative effects [39].

GC-MS analysis of ethanol leaf extract of *B. eurycoma* revealed the presence of 24 phytochemical compounds; Amongst which were the sugar moiety (3-O-Methyl-d-glucose and Ethyl .alpha.-d-glucopyranoside ), aldehyde (cis-9-Hexadecenal, Furan-3-carboxaldehyde-2-methoxy-2,3-dihydro-), Glucosinolates (Desulphosinigrin), Diterpene (phytol), phenol (Hydroquinone, 2,3-dihydro- Benzofuran), fatty acids esters (n-Hexadecanoic acid, 9,12-Octadecadienoic acid (Z,Z)-, Hexadecanoic acid, Hexadecanoic acid, 2-hydroxy-1-

(hydroxymethyl), Octadecanoic acid) and Flavonoid (4H-Pyran-4-one-2,3-dihydro-3,5-dihydroxy-6-methyl-).

Phytochemical and ethnobotanical database has ascribed several pharmacological or biological activities such as anti-inflammatory, antioxidant, antilipidemic, antihistaminic, antimicrobial amongst others to majority of the compounds identified [28]. 3-O-Methyl-d-glucose, the most abundant component in *B. eurycoma*, is a non-metabolizable glucose analog, used as a proxy for d-glucose uptake in *in vivo* absorption studies and as a preservative [40, 41].

Research findings have reported the anticancer and antimicrobial nature of desulphosinigrin [42]; antioxidant and antifungal activities of 4H-Pyran-4-one,2,3-,dihydro-3,5-dihydroxy-6-methyl- [43-45]; 5-Alpha reductase inhibitor, anti-androgenic, anticancer, antioxidant, hypocholesterolemic, acne reductive, anti-inflammatory and anti-eczemic effect of 9,12-Octadecadienoic acid (Z,Z)- [46-48], as well as the anti-inflammatory, anti-cancer, cytotoxic and antimicrobial properties of n-Hexadecanoic acid and Hexadecanoic acid [49-51].

Phytol, a building block of chlorophyll, is among the twenty four compounds identified in the present study. It is used in the manufacture of synthetic vitamins E and $K_1$. It has been reported to have antinociceptive, antioxidant, anti-inflammatory, antiallergic, immunostimulant, antimicrobial, antischistosoma, cancer preventive, sedative and anxiolytic effects [52-60]. Hydroquinone occurs naturally in various medicinal plants [61, 62], it has been reported to be allelochemic, antimicrobial, antihepatomic, antilithic, antimelanomic, antimelasmic,

uroantiseptic , antimitotic, and antipertussive [63-65]. Benzofuran, 2,3-dihydro is a coumaran and research has shown that it possesses anti-inflammatory, antidiarrheal, antileishmanial, immunomodulatory, antimicrobial and anti-helminthic activities [66-68]. Octadecanoic acid is a flavouring agent which is hypocholesterolemic [69]. 2,3-dihydro-3,5-dihydroxy-6-methyl-4H-Pyran-4-one, a flavonoid with antifungal, antioxidant [70, 71].

## Conclusion

This study showed that *B. eurycoma* leaves are nutritious and contains important phytochemicals with several biological activities revealing its potentials use in food, pharmaceutical, cosmetic and nutraceutical industry.

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