Isolation of a pentacyclic triterpenoid from the antiplasmodial bioactive fraction of *Nauclea latifolia* (Sm) roots

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Abstract

**Purpose:** To research the antiplasmodial property of aqueous extract, fractions, and residue of *Nauclea latifolia* roots and to isolate the components responsible for the antiplasmodial activity.

**Methods:** Roots of *N. latifolia* were macerated with distilled water; the extract was obtained, successively partitioned with ethyl acetate and butanol. The extract, fractions, and the residue obtained were evaluated for their in vivo antiplasmodial activity and compared with amodiaquine and artesunate. The residue (which exhibited the best therapeutic index) was subjected to column and thin layer chromatography to isolate its components. Purification led to the isolation of betulinic acid, which was characterized with the aid of spectroscopic techniques (¹H, ¹³C NMR, and EI-MS).

**Results:** The residue significantly inhibited parasite growth from 42.8 % (D2) to 77.6 % (D5). Therefore, residue exhibited the highest therapeutic index against Plasmodium berghei in the three in vivo antiplasmodial (prophylactic, suppressive, and curative) models and it compared favourably with amodiaquine (80.5 %) and artesunate (85.9 %). The major component of the residue was betulinic acid.

**Conclusion:** The results validate the antiplasmodial claims of the roots of *N. latifolia* in folkloric medicine, and demonstrated that the isolate has a high therapeutic index in this regard. Further investigations, however, are required to determine the clinical efficacy and safety of the compound/isolate.

**Keywords:** *Nauclea latifolia*, Malaria, Betulinic acid, Plasmodium berghei, Parasite density, Growth inhibition

INTRODUCTION

*Nauclea latifolia* (Rubiaceae) is also known as the Pincushion tree. It is highly distributed in the rainforest zone of Africa [1]. The stem, bark, root, leaf, and fruit are traditionally used by the local population as ethnomedicine for the treatment of pain, diabetes, infections, and malaria. Indole alkaloids with tetracyclic, or pentacyclic rings seem to be common in this family [2]. Moreover,
compounds such as naucleafine, naucleatine, nauclecholine, and naufoline [3,4], nauclefloline [5], strictosamide, naucleamides A, naucleamid F [6] have been isolated from the root bark of N. latifolia. Furthermore, five monoterpenic indole alkaloids, naucleamides A to E were isolated from the bark of the plant whereas the major indole alkaloid with amino acetal bridge and five cyclic ring structures was naucleamide E [7]. In addition, betulinic acid, a naturally occurring pentacyclic lupane-type triterpenoid, has been isolated from other plants and reported to have antiplasmodial activity [8,9].

A study involving thirty-three plants frequently used by trado-medical practitioners for the management of malaria in West Africa reported high antiplasmodial potentials of N. latifolia [10]. Consequently, in vitro antiplasmodial evaluation against P. falciparum revealed that the root extracts of N. latifolia were more potent than the stem extracts [1]. Preliminary studies on the plant showed that the aqueous extracts of N. latifolia roots exhibited a better antiplasmodial activity against strains of P. berghei as compared with the ethanol extract [11].

This prompted further work to isolate compounds from the bioactive fraction of the aqueous extract of N. latifolia roots.

**EXPERIMENTAL**

**Plant material**

The N. latifolia roots were obtained from Ikot Andem Ildidep, Ibiono Ibom L.G.A., Nigeria. Dr. Udo, a plant, a taxonomist in the Department of Pharmacognosy and Natural medicine, University of Uyo, Nigeria identified the roots whereas herbarium number UUH67G was documented. Thereafter, the roots were cut into smaller pieces, washed with running tap water, drained, dried, and pulverized. The aqueous extract was prepared by the maceration of the pulverized roots (5.2 kg) in boiled distilled water (5 L), kept for 72 h with occasional shaking, filtration was carried out using filter paper thereafter, the filtrate underwent concentration in a rotary evaporator at 40 °C. The repetition of the procedure yielded a more aqueous extract.

**Extraction and isolation**

The weight and % yield of aqueous extract was 246.9 g and 4.75 % respectively. Successive partitioning of the extract (150 g) with ethyl acetate (28x250 ml) and butanol (42 x 250 ml) using a separating funnel (1000 ml) yielded ethyl acetate fraction (13.66 g), butanol fraction (52.83 g), and residue (83.51 g) after the evaporation of the solvent. The extract, fractions, and residue were screened for acute toxicity and antiplasmodial activity. The residue was the most active against *Plasmodium berghei* in the three *in vivo* antiplasmodial models. The residue (10 g) was subjected to column chromatography (CC) using silica gel and eluted with the appropriate solvent systems (n-hexane, dichloromethane, ethyl acetate, and methanol) and yielded 358 fractions. These fractions were subjected to different phases of TLC analysis (pooling and bulking processes) and yielded twenty-four (24) and six (6) fractions (A-G) based on similarity in Rf values respectively. Bulked fraction C (3428 mg) was further subjected to CC using silica gel and eluted using appropriate solvent systems to yield 47 sub-fractions. These sub-fractions were identified with TLC to give six bulked sub-fractions (C1-C6). The sub-fraction C2 showed one spot on the TLC plate (using varying solvent systems) and was identified as betulinic acid (2205 mg). The sub-fraction C2 showed a single spot on TLC (using varying solvent systems) and was identified as betulinic acid (2205 mg).

**General procedures**

$^1$H and $^{13}$C NMR spectra were run on an Avance AV-400 MHz spectrometer. DMSO-$d_6$ enhanced the dissolution of sub-fraction C2 whereas trimethyl silane was utilized as referential ($\delta_H$, $\delta_C$= 0), with coupling constants in Hertz. EI-MS (positive mode) enhanced the quantification of mass spectrometry. Thin-layer chromatography was carried out by using Sigma–Aldrich’s analytical precoated TLC silica gel 60 and visualized under both long (366 nm) and short (254 nm) wavelength UV light. Moreover, Sigma–Aldrich’s silica gel Kiesel gel 60 (200–400 mesh, Merck) and Sephadex LH-20 enhanced column chromatography (CC) analysis.

**Phytochemical evaluation of Nauclea latifolia**

Standard procedures as described by Trease and Evans [12] were used to evaluate the qualitative phytochemical analysis of the extract, fractions, and residue of *N. latifolia* roots.

**Animals**

The Animal House in the Faculty of Pharmacy, University of Uyo, Nigeria provided one hundred and seventy-four (174) albino mice (13 – 27 g) used for the antiplasmodial experiment. The NIH protocols for the use, handling, and care of laboratory animals were judiciously followed [13]. The mice were housed at ambient temperature, they enjoyed access to water and feed (standard pellet diet) throughout the experiment. The
Animal Ethics Committee of the faculty approved the animal studies (approval no. UUFPHARM/0317) which followed international guidelines for animal studies.

Parasites

National Institute of Medical Research (NIMR), Nigeria provided the chloroquine-sensitive *P. berghei* (NK-65) infected donor mice that were used for the study.

Inoculum preparation

A stock of *Plasmodium berghei*-infected red blood cells with 20% minimal peripheral parasitemia was obtained by cardiac puncture of the parasitized mice and emptied into a tube containing anticoagulant. Evaluation of the number of *Plasmodium berghei*-infected erythrocytes against that of leucocytes enhanced the calculation of the percentage parasitemia. Normal saline was used for serial dilution and the final inoculum (0.2 ml) contained about 1x10^7 *Plasmodium berghei*-infected erythrocytes known as the standard inoculum for a mouse [14].

Drugs

Artesunate tablets (50 mg) from (Mekophar Chemical Pharmaceutical Joint-stock Company, Ho Chi Minh City-Vietnam) and amodiaquine tablets (200 mg) from (Pfizer Afrique de l’s Ouest Dakar R. P. Senegal) were dissolved in distilled water (100 ml) and given at dosages of 5 mg/kg and 30 mg/kg as positive controls in the antiplasmodial study.

LD50 determination of the plant extract

The LD50 and effective dosages of the plant’s root extracts were evaluated using Lorke’s method [15]. The aqueous extract, fractions, and residue (50 – 2000 mg/kg) were given through the peritoneum then, signs of toxicity signs such as gasping decreased respiratory rate, palpitation, and death were observed within 42 h (loss of consciousness, asphyxia, and death) were observed within forty-two (42) h post-administration.

Determination of *in vivo* antimalarial activity

The antiplasmodial potentials of the extract, fractions, and residue were determined by using curative, prophylactic, and suppressive models although with slight modifications of the experimental protocol as reported by Okokon et al [14].

Slide staining and examination

Blood from each mouse was collected and used in the preparation of two smears (thin and thick); nevertheless, there were slight modifications to the experimental protocol as earlier reported by Owusu-Agyei et al [16]. The Parasite densities/µL of the blood were counted against 200 or 500 leucocytes.

Statistical analysis

GraphPad Prism version 8.0 (GraphPad Software, Incorporated, San Diego, CA, USA) was used for data analysis, mean ± SEM is the format of result presentation whereas One-way ANOVA statistically analyzed the significant differences within and between the groups. Afterward, Turkey’s post hoc test enhanced multiple comparisons of means, and differences were adjudged statistically significant at a 95% confidence level.

RESULTS

Spectral characteristics

The spectral data of the isolated compound is presented as follows: the pure compound was observed as needle-like colorless substance; molecular weight and formula were given was 456 g/mol and C30H48O3 respectively; based on EI-MS, base peak and molecular ion peak at m/z as 189.1 and 456.4 respectively.

1H-NMR (DMSO, ppm): The details of proton spectrum (DMSO, δ ppm) are given as follows: the δH at 0.77 (s, 3-H, H-28), 0.88 (s, 3-H, H-29), 0.97 (s, 6-H, H-25 & 26), 1.02 (s, 3-H, H-30), 0.93 (s, 1-H, H-4), 1.06 (s,1-H, H-12), 1.42 (m, 8-H, H-6, 8,15 & 20), 1.58 (m, 4-H, H-7 & 14), 1.35 (m, 2-H, H-19), 1.71 (s, 3, H-14, 1.94 (m, 2H, H-16), 1.56 (m, 4-H, H1 & 13), 2.33 (m, 1H, H-21), 2.25 (m, 1H, H-18), 3.03 (m, 1H, H-2), 4.61 (s, 1-OH, H-27), 3.15 (dd, 1-H, H-23), and 3.03 (t, 1-H, H-23), representing different location and environment of proton present.

13C-NMR (DMSO, ppm): The detail 13C-NMR (DMSO, δ ppm) is as follows; 15.12 (C-30), 16.10 (C-28), 16.67 (C-7), 16.73 (C-29), 19.46 (C-14), 19.57 (C-24), 22.11 (C-25,26) 26.93 (C-13), 28.07 (C-1), 28.62 (C-15), 30.86 (C-20), 31.75 (C-16), 33.39 (C-8), 35.63 (C-19),38.16 (C-6), 39.70 (C-9), 39.96 (C-11), 48.58 (C-21), 49.29 (C-18), 50.51 (C-10), 56.91 (C-4), 57.53 (C-17), 79.70 (C-2), 110.3 (C-23), 152.03 (C-22), and 180.20 (C-31). For final confirmations, HMBC, HSQC, NOESY, DEPT-90, DEPT-135 and COSY techniques were used and the
structure of compound (betulinic acid) is presented in Figure 1.

![Chemical structure of betulinic acid](image)

**Figure 1:** Chemical structure of betulinic acid

**The prophylactic activity of extracts, fractions, and residue**

The repository antimalarial potentials of the extract, fractions, and residue of *Nauclea latifolia* roots were investigated and compared with amodiaquine, artesunate, and negative control groups for a test of significance at a 95% confidence level. Table 1 showed that the parasite densities in the entire group significantly reduced as compared with the negative control group. In ascending order of activity, *Plasmodium berghei*’s % growth inhibition was calculated as follows: butanol fraction (57.3 %) < residue (82.3 %) < artesunate (87.2 %) < ethyl acetate fraction (97.6 %) < amodiaquine (98.3 %) < aqueous extract (99.2 %). Moreover, the result of mean survival time (MST) revealed a significant elevation in all the groups except the butanol fraction that showed significant reduction as compared with the treatment group of amodiaquine.

**Table 1: Prophylactic activity of extracts, fractions, and residue**

| Treatment group                  | Mean weight (g) | Parasite density/µL | Mean survival time (days) |
|----------------------------------|-----------------|---------------------|---------------------------|
| Negative control (distilled water) | 17.5±0.3        | 49960.0±6491.6      | 14.5±0.3                  |
| Amodiaquine (30 mg/kg)           | 19.8±0.8        | 782.0±17.7a*        | 24.5±1.1 a**              |
| Artesunate (5 mg/kg)             | 18.5±1.3        | 6434.9±1167.5a*     | 19.8±1.3 a**              |
| Aqueous extract (150 mg/kg)      | 19.3±1.7        | 438.4±33.5a*        | 25.1±0.5 a**              |
| Ethyl acetate fraction (150 mg/kg)| 19.3±1.1        | 1276.3±59.1a*       | 22.4±1.7 a**              |
| Butanol fraction (150 mg/kg)     | 19.0±1.3        | 21398.0±867.9a,b*,c** | 16.7±0.6b*               |
| Residue (150 mg/kg)              | 18.3±0.5        | 8781.0±406.2a*      | 19.4±1.6 a**              |

*a* implies significant reduction at 95 % confidence level, the comparison was done with negative control whereas n = 6. b** illustrates significant elevation at 95 % confidence level, the comparison was done with amodiaquine whereas n = 6. c** implies significant elevation at 95 % confidence level, the comparison was done with artesunate whereas n = 6

**Suppressive activity of extracts, fractions, drugs, and residue**

The investigation evaluated mean survival time, parasite densities, and % chemosuppressive potentials of the extract, fractions, and residue of *Nauclea latifolia* roots, artesunate, and amodiaquine. Also, Table 2 indicated a significant reduction in parasite densities at a 95 % confidence level in the entire group as compared with the negative control. In descending order of activity, the % growth inhibition of *P. berghei* in this study was calculated as follows: amodiaquine (99.1 %) > artesunate (98.9 %) > aqueous extract (98.1 %) > residue (91.4 %) > butanol fraction (70.3 %) > ethyl acetate fraction (52.4 %). Moreover, the significant elevation in MST at a 95 % confidence level was recorded in the entire group as compared with the negative control group.

**Schizonticidal activity of extracts, fractions, and residue**

This study evaluated parasite densities, mean survival time, % growth inhibition of *Plasmodium berghei* to unravel the schizonticidal potentials of the infused extract, fractions, and residues of *Nauclea latifolia* roots at dosages (150, 300, and 450 mg/kg) and compared with the groups of negative control, amodiaquine, and artesunate. Table 3 indicated that there was a significant reduction in parasite densities at a 95 % confidence level in all the groups on day two (D2) and day five (D5) as compared with their respective negative control group. Also, significant elevation in MST at a 95 % confidence level was observed in all the treatment groups except ethyl acetate fraction as compared with the group of the negative control.
### Table 2: Suppressive activity of extract, fractions, and residue

| Treatment groups                        | Mean weight (g) | Parasite densities/µL | Mean survival time (days) |
|-----------------------------------------|-----------------|-----------------------|---------------------------|
| Negative control (distilled water)       | 15.5±0.7        | 47584.1±12361.0       | 12.2±1.5                  |
| Amodiaquine (30 mg/kg)                  | 18.3±0.9        | 390.7±93.4**          | 25.7±1.8*                 |
| Artesunate (5 mg/kg)                    | 16.3±0.8        | 560.1±105.6**         | 24.4±0.7**                |
| Aqueous extract (150 mg/kg)             | 18.0±1.5        | 978.1±239.7**         | 23.8±1.8**                |
| Ethyl acetate fraction (150 mg/kg)      | 16.8±1.3        | 22681.2±3647.1**      | 15.2±0.4**                |
| Butanol fraction (150 mg/kg)            | 18.3±0.8        | 14108.1±4198.2**      | 17.1±1.2                  |
| Residue (150 mg/kg)                     | 13.8±4.1        | 4119.1±383.2**        | 21.1±2.1**                |

*a* implies significant reduction at 95 % confidence level, the comparison was done with negative control whereas n = 6. *b* implies significant reduction at 95 % confidence level, the comparison was done with amodiaquine whereas n = 6. *c* implies significant reduction at 95 % confidence level, the comparison was done with artesunate whereas n = 6. 

### Table 3: Schizonticidal activity of extracts, fractions, and residue

| Treatment group                        | Parasite density/µL on D₀ & % growth inhibition | Parasite density/µL on D₂ & % growth inhibition | Parasite density/µL on D₅ & % growth inhibition | Mean survival time (days) |
|-----------------------------------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|---------------------------|
| Negative control (distilled water)      | 134021.0± 15652.0 (-32.2%)                     | 177052.0± 28312.0 (-66.4%)                    | 222876.1± 30656.1 (-80.8%)                    | 9.6± 0.24                 |
| Amodiaquine (30 mg/kg)                  | 130995.1± 16454.1 (52.2%)                      | 62746.0± 6998.0 (85.7%)                       | 25516.0± 2292.0 (80.8%)                      | 22.6±0.23**               |
| Artesunate (5 mg/kg)                    | 169896.1± 12742.1 (47.1%)                      | 90092.0± 3098.0 (85.7%)                       | 24004.1± 3745.1 (85.7%)                      | 24.2±1.33**               |
| Aqueous extract (150 mg/kg)             | 204132.1± 10805.1 (3.4%)                       | 99256.0± 2273.0 (51.4%)                       | 57679.1± 494.6 (71.7%)                       | 19.66±0.57**              |
| Aqueous extract (300 mg/kg)             | 241429.1± 40596 (63.4%)                         | 88112.0± 14842.0 (77.8%)                      | 54017.2± 17178.1 (77.8%)                     | 22.2±0.43**               |
| Aqueous extract (450 mg/kg)             | 236963.1± 51136.1 (49.9%)                      | 118903.0± 13298.0 (74.6%)                     | 60522.0± 14058.1 (74.6%)                     | 20.4±0.36**               |
| Ethyl acetate fraction (300 mg/kg)      | 168675.1± 18613.1 (13.7%)                       | 145654.0± 24429.0b (47.2%)                    | 89443.2± 9388.1a* (47.2%)                    | 15.3±0.36b*, c*           |
| Butanol fraction (300 mg/kg)            | 171646.1± 8238.6 (30.3%)                       | 120106.0± 21748.0 (64.2%)                     | 61678.1± 11616.1a* (64.2%)                   | 17.1±0.42a**, c*          |
| Residue (300 mg/kg)                     | 206217.1± 43858.1 (42.9%)                      | 117922.0± 16843.0 (77.8%)                     | 46138.1±5267.0a* (77.8%)                    | 22.2±0.46a**              |

** implies elevation at 95 % confidence level, the comparison was done with baseline parasite density on day 1 (D₀). *a* implies significant reduction at 95 % confidence level, the comparison was done with negative control whereas n = 6. *b** implies significant elevation at 95 % confidence level, the comparison was done with negative control whereas n = 6. *b* implies significant reduction at 95 % confidence level, the comparison was done with amodiaquine whereas n = 6. *c* implies significant reduction at 95 % confidence level, the comparison was done with artesunate whereas n = 6.
Table 4: Phytochemical screening of the extract, fractions, and residues of Nauclea latifolia roots

| S/no. | Test                          | Infused aqueous extract | Ethyl acetate fraction | Butanol fraction | Residue |
|-------|-------------------------------|-------------------------|------------------------|------------------|---------|
| 1.    | Alkaloids                     |                         |                        |                  |         |
|       | Dragendorff’s reagent         | +++                     | +                      | ++               | +++     |
|       | Hagger’s reagent              | +++                     | +                      | ++               | +++     |
| 2.    | Saponins                      |                         |                        |                  |         |
|       | Froth test                    | +++                     | ++                     | +++              | ++      |
|       | Emulsion test                 | +++                     | ++                     | ++               | ++      |
|       | Fehling’s test                | +++                     | ++                     | ++               | ++      |
| 3.    | Tannins                       |                         |                        |                  |         |
|       | Ferric chloride               | +                       | +++                    | +++              | +       |
| 4.    | Flavonoids                    |                         |                        |                  |         |
|       | Magnesium metal               | ++                      | ++                     | ++               | ++      |
|       | Ethyl acetate                 | ++                      | ++                     | ++               | ++      |
| 5.    | Glycoside                     |                         |                        |                  |         |
|       | Hydrolysis                    | ++                      | +                      | ++               | ++      |
| 6.    | Cardiac glycoside             |                         |                        |                  |         |
|       | Salkowski test                | +++                     | +++                    | ++               | ++      |
| 7.    | Terpenes & steroids           |                         |                        |                  |         |
|       | Liberman-Burchard             | ++                      | +                      | +                | ++      |
| 8.    | Deoxysugar                    |                         |                        |                  |         |
|       | Keller-Killiani test          | ++                      | +                      | +++              | ++      |
| 9.    | Anthraquinone                 | _                       | _                      | _                | _       |
| 10.   | Carbohydrate                  | ++                      | ++                     | +++              | +++     |

+++ implies high relative abundance, ++ stands for moderate relative abundance, + connotes low relative abundance, and - implies absence.

The qualitative phytochemical evaluation of extract, fractions, and residue of plant roots is presented below.

**DISCUSSION**

Some secondary metabolites like alkaloids, terpenoids have been reported to have antiplasmodial activity [17] therefore, the abundance of alkaloids, saponins, carbohydrates, terpenes, cardiac glycosides, and flavonoids in the extracts, fractions, and residues of *N. latifolia* roots corroborated the assertion by these researchers hence, its traditional usage in the treatment of malaria in Nigeria.

Also worthy of note is the median LD$_{50}$ (1500.19 mg/kg body weight of mice) of the plant extract deduced as the geometrical mean of the highest dosage that produced zero % death in mice. The value revealed medial lethality of the plant extract with the possibility of inducing minimal multi-organ toxicity when consumed as a concoction or as an herbal formulation.

Moreover, the significant reduction in the parasite density in the entire groups as compared with the group of negative control revealed the varying degrees of repository activity of the plant’s extract, fractions and residue hence, the consumption of the root *N. latifolia* infusion may avert malaria incidence as well as its symptoms through the destruction of the pre-erythrocytic stage of the parasite. Furthermore, percentage growth inhibition of parasites is one of the indices when evaluating drug efficacy in anti-plasmodial studies so, the higher the value the better the therapeutic index of the drug. The result of the repository activity of the plant’s extract, fractions, and the residue was in tandem with the reports by Boampong et al [17] on the prophylactic efficacy of the extract of *Haematostaphis barteri* stem bark and 49.58% for ethanol extract of *N. latifolia* stem bark at the dosages of 300 mg/kg [18,19]. The corroboration of parasite density reduction, increased percentage growth inhibition of parasite with increased mean survival time of the mice suggests that the root *N. latifolia* has good repository activity.

The study further showed that the plant roots had good chemosuppressive potentials due to their...
improved parasite density reduction, increased mean survival time, and percentage growth inhibition of Plasmodium berghei. Whereas some researchers reported the inhibition of protein synthesis in parasites [20, 21] and the possibility of alkaloids blocking protein synthesis in P. falciparum [22] as the possible mechanism of actions of some plant extracts. So, the plant roots may have acted through a similar mechanism of action due to the abundance of alkaloids in them. In addition, the study showed higher percentage growth inhibition of the parasite when compared with 55.61% of aqueous extract of Parkia biglobosa administered at a dosage of 600 mg/kg [23] as well as 82.18% and 61.87% for Clerodendrum myricoides leaves and roots respectively [24]. Interestingly, the chemosuppressive activity of the residue group compared favorably with the artesunate and amodiaquine drugs, due to the presence of betulinic acid, a compound already reported to have anti-plasmodial activity. Therefore, the similarity in therapeutic index of this plant’s roots with the conventional drugs showed that the plant has good chemosuppressive potentials against P. berghei [18], it might use a similar mechanism of action as artesunate and amodiaquine drugs as well as being a candidate in the quest of profiling and designing drugs to avert high mortality rate from malaria, aggravated by the prevalence of drug-resistant strains of Plasmodium falciparum.

Since anti-plasmodial microscopic investigation targets the schizont stage of the parasite as the stage that causes clinical symptoms in patients; so, its evaluation gives clue on the therapeutic efficacy of a drug. Therefore, the schizonticidal activity of the extracts, fractions, and residue investigated in this study showed that the significant reduction in their parasite densities as compared with the negative control group was consistent with similar studies like 75.4% for the aqueous extract of N. latifolia stem bark [25] and 81-91% for the extract of N. pobeguini [26]. Hence, the validated schizonticidal activity of these plant roots may arise due to betulinic acid alone or its interactions with other bioactive compounds by directly inducing cytotoxicity on P. berghei. In addition, the spectral data from 13C-NMR (DEPT-135, BB, and DEPT-90) revealed the compound has 30 carbons atoms constituting six methines, six methyls, seven quaternary carbon, and eleven methylenes atoms. Also detected were five rings, one double bond, and one acidic carbonyl in the molecule. 1H-NMR revealed six singlet methyls, a carbinolic proton, a pair of olefinic protons with an exocyclic methylene group which was characteristic for lupane triterpenes [27]. Furthermore, the fragmentation pattern from EI-MS was typical of lupane triterpenes through the presence of m/z 438 [M-H2O], 411 [M-COOH], 248 [C16H24O2], 220 [C14H20O2], 205 and 207 [M - C15H27], 203 [248 - COOH], 203 [220 - OH], 189 [207 - H2O], 175 [220 - COOH]. Based on the foregoing spectral data, the compound was similar to those earlier reported for betulinic acid [27].

CONCLUSION

The plant root extract of N. latifolia and its fractions have been bioactively screened via in vivo anti-plasmodial studies while the residue with the best therapeutic indices have been subjected to column chromatography, and the eluent NLA1 characterized with the aid of spectroscopic techniques. The isolation of betulinic acid from the Nauclea genus is reported here for the first time although the pentacyclic triterpenoid had previously been reported to possess anti-plasmodial activities in other plants. Hence, the high therapeutic indices observed in this study may solely be due to betulinic acid or its synergistic interaction with other compounds. Further investigations, however, are required to determine the clinical efficacy and safety of the compound/isolate.

DECLARATIONS

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Conflict of interest

No conflict of interest is associated with this work.

Contribution of authors

We declare that this work was done by the authors named in this article and all liabilities about claims relating to the content of this article will be borne by the authors. Asanga, Edet Effiong conceived, designed the study; collected and analyzed the data as well as wrote the manuscript. Igile, Godwin O., Ebong, Patrick E., and Eseyin, Olorunfemi A. read and approved the manuscript. Emmanuel E. Essien and Thomas Paul Sunday collected and analyzed the data.
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