Research Article

Anti-Onchocercal Properties of Extracts of *Scoparia dulcis* and *Cyclicodiscus gabunensis*

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Introduction. The elimination of onchocerciasis is hampered by the absence of suitable drugs that are effective against adult filariae. This study is aimed at assessing the anti-onchocercal effects of extracts of *Scoparia dulcis* and *Cyclicodiscus gabunensis* that could serve as drug leads against onchocerciasis. Methods. Different parts of the plants (*Scoparia dulcis* and *Cyclicodiscus gabunensis*) were extracted with hexane, methylene chloride, and methanol. The extracts were tested in vitro against the bovine model parasite, *Onchocerca ochengi*. Adult female worm viability was determined biochemically by MTT/formazan colorimetry, while the adult male and microfilariae viability were determined by microscopy based on % inhibition of worm motility score. Cytotoxicity and acute toxicity of active extracts were tested on monkey kidney epithelial cells (LLC-MK2) and Balb/C mice, respectively. Results. The hexane extract of *Scoparia dulcis* recorded the highest activity, with IC50s of 50.78 μg/ml on both adult male and female worms and 3.91 μg/ml on microfilariae. For *Cyclicodiscus gabunensis* extract, the highest activity was seen with the methylene chloride extract, with IC50s of 50.78 μg/ml, 62.50 μg/ml, and 16.28 μg/ml on, respectively, adult male, female, and microfilariae. The 50% cytotoxic concentration on the LLC-MK2 cells was 31.25 μg/ml for the most active extracts. No acute toxicity was recorded for the extracts. Phytochemical analysis of the extracts revealed the presence of alkaloids, flavonoids, sterols, saponins, phenols, and glycosides. Conclusion. This study validates the traditional use of these plants in treating onchocerciasis and suggests *S. dulcis* and *C. gabunensis* as new potential sources for the isolation of anti-onchocerca lead compounds.

1. Introduction

Medicinal plants are widely used in traditional cultures all over the world and are becoming increasingly popular in modern society to meet health care needs, particularly in Africa and most developing countries [1]. Natural product molecules represented more than 50% of drugs that had been put into the drug market, with many medicinal plants extracts now used as prescription drugs in many developed countries [2–4]. Approximately, a quarter of all Food and Drug Administration and European Medical Agency (EMA) approved drugs are plant-based, with well-known drugs such as quinine, artemisinin, morphine, and the anticancer drugs paclitaxel and vincristine [5–8]. The presence of phytoconstituents gives plant species their medicinal potential with numerous functions such as antimicrobial, anticancer, antioxidant, and antiviral [1, 9]. *Scoparia dulcis* is a herb of the family Plantaginaceae that is common in Tropical Africa, Asia, and Central America. It is used traditionally to treat fever, cold, sore throat, and eczema, with other ethnomedical uses like the treatment of gastric problems, reproductive issues, piles, liver, and respiratory diseases [10, 11]. Approximately, 160 compounds have been identified from *S. dulcis*, among which 155 have been related to the treatment of metabolic syndromes [10]. Previous chemical analysis of the plant has identified various phytoconstituents such as nitrogen-containing compounds, flavonoids, diterpenoids, triterpenoids, steroids, phenolics,
and aliphatics, with several pharmacological effects such as been antidiabetics, anticancer, antiarthritic, anti-hyperlipidaemia, anti-inflammatory, and antiuriculithiasis [10, 12, 13].

_Cylicodiscus gabunensis_ (Fabaceae) is an indigenous medicinal plant that is widely distributed in West and Central Africa. Extracts from its stem bark have been used for the treatment of viginitis, jaundice, amalaria, antibacteria, anti-inflammatory, antipyretics, and as soaps and mouthwash by some communities [14–17]. Previous phytochemical analysis has revealed the presence of triterpenoids, saponins, and phenolics with some definite pharmacological antimalarial, antifree radical, and antibacteria properties [16, 18, 19]. Ethnopharmacological information about _S. dulcis_ and _C. gabunensis_ indicated their use by an ethnic group in Cameroon to treat filarial infections like onchocerciasis.

Human onchocerciasis, or river blindness, is a neglected tropical disease with serious debilitating effects, affecting over 37 million people worldwide, mostly in developing countries, including Yemen and many countries found in Africa, Central and South America [20]. Over 200 million people are at risk of onchocerciasis infection, and more than 99% of the disease burden is found in Africa, with an estimated 1.15 million of the infected being blind and an additional 500,000 living with severe visual impairment [21–23]. The disease is caused by the filarial parasitic nematode _Onchocerca volvulus_, with the Simulium blackfly being its vector. Despite several efforts to control the disease, limitations are found in new strategies for blocking transmission [24]. Ivermectin remains the only drug currently recommended for the treatment and control of the disease. The drug mainly targets the microfilarial (juvenile) stage of the parasite, leaving the adult worms to continue to reproduce. There is also a serious adverse effect in individuals who are treated with ivermectin and are coinfected with _Loa loa_ with high microfilaraemia (greater than 30 000 microfilariae per ml) [25]. Reports on the resistance of the parasites to some of the filaricidal drugs, coupled with some of their drawbacks and restrictions of use in some endemic areas, have led to the urgent search for alternative treatments with macrofilaricidal properties [7, 25]. One strategy employed has been the exploitation of medicinal plants for the identification of novel potential drug leads. This study investigated the filaricidal properties of _Scoparia dulcis_ (Plantaginaceae) and _Cylicodiscus gabunensis_ (Fabaceae) on cattle derived from _Onchocerca ochengi_ (the closest known relative of _Onchocerca volvulus_), and their possible use as sources of new drug leads for the treatment of onchocerciasis.

2. Materials and Methods

2.1. Ethics Approval and Consent to Participate. Ethical clearance (2019/018/UB/IACUC/BU/TU/FS) for the use of the animals was obtained from the Institutional Animal Care and Use Committee (UB-IACUC), University of Buea, Cameroon.

2.2. Collection and Identification of Plant Materials. Aerial parts of _Scoparia dulcis_ and stem bark of _Cylicodiscus gabunensis_ were collected from Okpambe village in the Takamanda area in the Manyu Division South West Region of Cameroon in March 2016. The plants were selected based on ethnopharmacological information about them. They were identified and authenticated at the Limbe Botanic Garden, South West Region of Cameroon, and given the voucher numbers, SCA 1225 and SCA 5092, for _Scoparia dulcis_ and _Cylicodiscus gabunensis_, respectively.

2.3. Preparation of Plant Crude Extracts. The harvested plants were air-dried and ground to a fine powder. The ground material was weighed and subsequently immersed and macerated for 72 hours in three different solvents, namely: hexane (Hex), methylene chloride (MeC), and methanol (MeOH). For each solvent, the maceration was repeated twice. The mixture was filtered and the filtrate was concentrated using a rotary evaporator (BUCHI Rotavapor R-200, Switzerland) at appropriate temperatures. The concentrates were recovered with methylene chloride and allowed to stand at room temperature until the residual solvents had evaporated. The dried crude extracts were stored at −20°C until needed for assays. Stock solutions of 25 mg/ml of the different plant extracts were prepared in >99.8% dimethyl sulfoxide (DMSO) (Sigma, USA) and stored at −20°C until tested in biological assays.

2.4. Screening against _O. ochengi_ Adult Worms. The adult worms were extracted according to Cho-Ngwa et al. 2010. Briefly, subcutaneous nodules containing adult _O. ochengi_ worms were identified on the umbilical skin of infected cows and immediately brought to the laboratory. Under sterile conditions, adult worm masses containing one viable adult female and zero to several adult males were carefully recovered by dissection of the nodule with a sterile razor blade. The masses were then incubated in 2 ml of complete culture medium (CCM), which comprised RPMI-1640 (Sigma-Aldrich, U.S.A), 10% newborn calf serum, 200 units/ml penicillin, 200 μg/ml streptomycin, and 2.5 μg/ml amphotericin B (Sigma-Aldrich, U.S.A) in standard 12-well culture plates (NUNC, USA) and incubated overnight at 37°C in a 5% CO₂ incubator (Thermo Fisher, UK). The next day, 1000 μg/ml of the different plant extracts was prepared in 2 ml of CCM to generate a final of 500 μg/ml for the primary screening. The extracts and controls were tested in triplicate at each concentration, and the experiment was repeated thrice on different days. The negative control wells contained only the diluent, DMSO (500 μg/ml). Cultures were terminated on day 7, post addition of plant extracts. Adult male worm viability was visually scored using an inverted microscope (Nikon Eclipse TS200, China) on days 5 and 7, with a percentage reduction of motility ranging from 100% (complete inhibition of motility), 75% (only head or tail of worm motility), 50% (whole body of worm motile but sluggishly), 25% (almost vigorous motility) to 0% (no observable reduction in motility) [26]. Adult female worm viability was assessed on day 7 by the standard MTT/
Formazan assay in which each nodular mass was placed in a well of a 48-well microtitre plate containing 500 μL of 0.5 mg/ml MTT (Sigma-Aldrich, U.S.A) in incomplete culture medium (ICM) (composed of RPMI-1640 (Sigma-Aldrich, U.S.A), 200 units/ml penicillin, 200 μg/ml streptomycin, and 2.5 μg/ml amphotericin B, and then incubated in the dark at 37°C for 30 minutes. Adult female worm viability was evaluated visually by the extent to which the female worm mass was stained with MTT. Mean percent inhibition of formazan formation was calculated relative to the negative control worm mass. Adult worm death is positively correlated with the inhibition of formazan formation. The positive control was auranofin at 30 μM [25]. Secondary screening for the active extracts (100% activity) was done to determine the IC₅₀ for the extracts. The active extracts were restested as described under the primary screen at serial dilutions of seven concentrations (from 500 to 7.8125 μg/ml). Assays were done in triplicate and each experiment was repeated for confirmation. The means of all activities at a concentration were calculated using the statistical analysis Graphpad Prism version 6.0 (Graphpad Software, CA, USA) to generate dose-response curves from which the IC₅₀ values were obtained.

2.5. Screening against O. ochengi Microfilariae in Vitro. The mammalian kidney cells (LLC-MK2) obtained from the American Type Culture Collection (ATCC, Virginia, USA) were proliferated in CCM at 37°C under an atmosphere of 5% CO₂ in humidified air. The cells were seeded in 96-well plates until they became fully confluent and served as feeder layers for the mf assays. The cells were also used for cytotoxicity assessment of the extract [27]. O. ochengi microfilariae were isolated by the method of Cho-Ngwa et al. [28] with slight modifications. Briefly, umbilical cattle skin pieces containing palpable nodules were obtained from the abattoir, cleaned, carefully shaved, and sterilized with 70% ethanol. Skin slivers were obtained and incubated for 4–6 hours at room temperature in CCM. The emerged O. ochengi mf was concentrated by centrifugation (400xg, 10 min.). The highly motile mf was resuspended in CCM and distributed into wells (approximately 15 μl/100 μl of CCM/well) of 96-well plates containing the LLC-MK2 cell layer, and their viability and sterility were ascertained at the 24th hour prior to the addition of extracts.

The primary screens for O. ochengi mf were done at 500 μg/ml in duplicates to eliminate inactive extracts. Extracts that showed 100% activity were restested in the secondary screening as described above for the adult worm assay to determine the IC₅₀ values. The mf was incubated at 37°C under an atmosphere of 5% CO₂ in humidified air for 5 days. The positive control was amocarazine (5 mM), while the negative control contained 2% DMSO. Mf motility was scored microscopically daily. The percentage motility inhibition scores were assigned as 100% (all mf immotile), 75% (only head or tail of mf shaking occasionally), 50% (whole body of mf motile but sluggishly or with difficulty), 25% (almost vigorous motility), and 0% (vigorous motility) [28, 29].

2.6. Cytotoxicity Studies. This experiment was conducted on the active extracts only. The fully confluent cells were cultured in the presence of the extracts at varying concentrations and observed daily under an inverted microscope. By day 7, an MTT colorimetric assay was done on the cells [30]. Succinctly, MTT colorimetric assay involved the culturing of LLC-MK2 cells in CCM in a 96-well flat bottom plate, until when the cells attain a density of approximately, 3000 cells/100 μl of CCM. Thereafter, the culture media of the fully confluent cells was removed by inverting, flicking, and blotting the plate, followed by the addition of crude extracts in fresh CCM. Colored extracts were washed twice with 100 μl of ICM. A 5 mg/ml MTT stock in 1x phosphate buffer saline was prepared and diluted in ICM to obtain a concentration of 1 mg/ml. To the cells in each well was added 100 μl of the 1 mg/ml solution of MTT. The plate was incubated for 3 hours at 37°C in a humidified 5% CO₂ incubator. At the end of incubation, the MTT solution was removed by inverted flicking and blotting of the plate. One hundred microliters (100 μl) of DMSO was added to each well and the plate was shaken at 400 rpm for 5 minutes. Inhibition of formazan formation with MTT directly correlates with cell death, purple formazan formation indicated viable cells, while pale purple correlated with the IC₅₀ of the cells [30].

Under an inverted microscope (Nikon Eclipse TS200, China), living cells are flattened out and attached to the culture plate, while dead cells are rounded up and detached from the bottom of the plate.

The selectivity index (SI) values were calculated using the ratio as follows:

\[ SI = \frac{IC₅₀ of mammalian cells}{IC₅₀ of parasite} \]

2.7. Acute Toxicity Studies. The test was conducted in accordance with the OECD guidelines for testing of chemicals [31] and the animal protocol was approved by The Animal Care and Use Committee, of the Faculty of Science, University of Buea. The extracts with antifilaricidal activity were tested for acute toxicity in BALB/c mice. A total of 12 animals of approximately 20 g of body weight each were used and divided into 2 groups, 6 for each extract and of equal sex. Each of the 2 treatment groups received one of the extracts at a limited dose of 2000 mg/kg body weight, administered orally in a maximum volume of 250 μl of vegetable oil per animal, while the control group received the diluent only. The animals were observed daily for 14 days for any changes in physical activity, food intake, and water intake, loss of fur, stool samples, sensitivity to sound, and sensitivity to pain, motility, and mortality.

2.8. Phytochemical Analysis. The phytochemical properties of the active extracts were determined qualitatively using standard procedures: frothing test for saponins, Fehling’s test for glycosides, Meyer’s and Dragendorff’s reagents for alkaloids, acetic anhydride and sulphuric acid reagents for
steroids, ferric chloride test for tannins, aluminium chloride test for flavinoids, and ferric chloride and alcohol test for phenols [32].

2.9. Statistical Analysis. Data were analyzed using Graphpad Prism version 6 (Graphpad Software, CA, USA) to determine the IC50s and mean motility scores.

3. Results and Discussion

A total of 6 crude extracts were prepared from the two plants using solvents of different polarities (hexane, methylene chloride, and methanol). Results of the primary screen showed that the hexane and methylene chloride extracts of the two plants were more effective in killing the adult *Onchocerca ochengi* male, female, and microfilaria with IC50 values within the range of 62.5–3.91 μg/ml (Table 1). The hexane extract of *Scoparia dulcis* was more active against adult male and female worms, with an IC50 value of 50.78 μg/ml, while the methylene chloride extract was active with IC50 values of 62.50 μg/ml and 60.57 μg/ml on male and female *O. ochengi*, respectively. Both extracts exhibited the same activity on microfilaria, with an IC50 value of 3.91 μg/ml (Figures 1(a)–1(c)). The methylene chloride extract of *Cyclodiscus gabunensis* were active against adult male, female, and microfilaria of *O. ochengi* with IC50 values of 50.78 μg/ml, 62.50 μg/ml, and 16.28 μg/ml, respectively, while the hexane extract recorded an IC50 value of 22.10 μg/ml for the *O. ochengi* microfilaria (Figures 1(a)–1(c)).

The time-dependent studies of SDHex and SDMeC demonstrated a 100% inhibition of mf motility on the 72nd hour for both extracts at 7.81 μg/ml, while those of CGHex and CGMeC showed 75% and 100% inhibition, respectively, for mf motility on the 120th hour at 31.25 μg/ml (Figure 2).

<table>
<thead>
<tr>
<th>Extracts Code</th>
<th>IC50 (μg/ml) for % inhibition of male</th>
<th>IC50 (μg/ml) for % inhibition of male</th>
<th>IC50 (μg/ml) for % inhibition of female</th>
<th>IC50 (μg/ml) for % inhibition of female</th>
<th>IC50 (μg/ml) for % inhibition of microfilaria</th>
<th>IC50 (μg/ml) for % inhibition of microfilaria</th>
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<tr>
<td>SDHex</td>
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<td>62.50</td>
<td>50.78</td>
<td>62.50</td>
<td>3.91</td>
<td>7.81</td>
</tr>
<tr>
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<td>60.57</td>
<td>125</td>
<td>3.91</td>
<td>7.81</td>
</tr>
<tr>
<td>CGMeC</td>
<td>50.78</td>
<td>62.5</td>
<td>62.5</td>
<td>125</td>
<td>16.28</td>
<td>31.25</td>
</tr>
<tr>
<td>CGHex</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>22.10</td>
<td>62.50</td>
</tr>
</tbody>
</table>

(SDHex: hexane extract of *S. dulcis*; SDMeC: methylene chloride extract of *S. dulcis*; CGMeC: methylene chloride extract of *C. gabunensis*; CGHex: hexane extract of *C. gabunensis*).

The active extracts were selected for acute toxicity studies in BALB/C mice at a limited dose of 2000 mg/kg of body weight. Following the administration of the test substance to that of the control mice, no change was observed in the physical activity and behavior of the test and control animals. The test and control groups were indistinguishable from one another on the basis of their appearance and physical activity at the end of the 14 days study period.

3.2. Phytochemical Composition of Active Extracts. Phytochemical screening revealed the different classes of secondary metabolites present in the active hexane and methylene chloride extracts of *S. dulcis* and *C. gabunensis*. The *S. dulcis* extracts showed positive for the presence of steroids, glycosides, phenolics, and flavonoids (Table 3), with alkaloids and saponins present only in the hexane and methylene chloride extracts, respectively. The *C. gabunensis* extracts showed positive for the presence of steroids, glycosides, phenolics, and flavonoids (Table 3), with alkaloids and saponins present only in the hexane and methylene chloride extracts, respectively.

4. Discussion

The aim of this study was to appraise the anti-onchocercal activity of two medicinal plants—*Scoparia dulcis* and *Cyclodiscus gabunensis*. From the ethnopharmacological survey, the plants were selected for the study based on their traditional claim to manage filariasis by traditional health practitioners in Okpambe; a village in the Takamanda area in the Manyu Division of the South West Region Cameroon. There has not been any scientific report on the use of these plants and its metabolites for the management of filaria infections. Previous studies of the plants have been reported for their use as antidiabetes, anti-inflammatory, antimicrobial, antisickling, antioxidant, antimalarial, and in the treatment of skin diseases, warts, jaundice, dysentery, gonorrhea, snake bites, and gastrointestinal disorders [11, 15, 33, 34]. *Onchocerca ochengi* was considered the most suitable model for screening anti-onchocerca phytomedicines because it is the closest relative to *Onchocerca volvulus* in which both parasite shares the same Simulium vector, in addition to the low cost of obtaining *O. ochengi* and the ease of its availability [25, 26].

The primary screen was done *in vitro* on all stages of the *Onchocerca ochengi* parasite for the hexane, methylene chloride, and methanol extracts of *S. dulcis* and
C. gabunensis plants. The hexane and methylene chloride extracts showed 100% activity at 500 μg/ml against the stages of the parasite. The extracts were subjected to a secondary screen at lower doses, ranging from 500 μg/ml down to 7.8125 μg/ml. This gave promising IC50 values within the range 62.5–3.91 μg/ml. The hexane extract of S. dulcis was active against the adult male and female O. ochengi with IC50 value of 50.78 μg/ml, while the methylene chloride extract showed activity with IC50 values of 62.50 μg/ml and 60.57 μg/ml on the male and female O. ochengi, respectively. The methylene chloride extract of C. gabunensis was active against the adult male and female of the parasite, with IC50 values of 50.78 μg/ml and 62.50 μg/ml respectively, (Figures 1(a) and 1(b)). The activity against the adult onchocerca worms is an important finding because as of date, the lone approved recommended drug for the treatment of the disease does not kill the adult worms (macrofilariae), leaving it to continue to reproduce and produces the microfilariae (juvenile worms) that generate pathologies [25]. The hexane and methylene chloride extracts of S. dulcis exhibited the same activity on the microfilariae with an IC50 value of 3.91 μg/ml, while the hexane extract of C. gabunensis recorded an IC50 value of 22.10 μg/ml (Figure 1(c)). These results add information in support of previous findings that showed nonpolar extracts to be nematocidal more than polar ones [27, 35, 36]. The seemingly same IC50 values of both extracts on the male and female Onchocerca parasites might be due to the fact that both parasites have the same targets.

**Figure 1:** (a) % inhibition of O. ochengi adult male motility for SDHex, SDMeC and CGMeC (b) % inhibition of formazan formation on O. ochengi adult female motility for SDHex, SDMeC and CGMeC (c) % inhibition of O. ochengi mf motility.
for the active plant extracts, while the discrepancy in activity between the adult and juvenile parasites probably indicates a variation in the target of the different extracts. The juvenile form of the parasites was more liable to be killed than the adult worms. This might be due to the fact that the juvenile worms are not entangled in nodular masses like adult worms, giving them more exposure to the compounds than adult parasites [29]. In addition, the fact that the extracts could kill the different parasitic stages of *O. ochengi* making these plants a potential source for the isolation of novel compounds with anti-onchocerca activity.

The safety of the plant extracts was done by evaluating cytotoxicity in LLC-MK2 cells and acute toxicity in Balb/c mice. The selectivity index of *S. dulcis* extracts was less than 1 for the adult parasites and 7.99 (greater than 1) for the mf, while that of the *C. gabunensis* extracts was all less than 1. These values indicate that the hexane and methylene chloride extracts of *S. dulcis* were more toxic to the juvenile parasites than the LLC-MK2 cells. The high toxicity of the extracts on the adult parasites might be due to the complex morphology of the adult parasites relative to the LLC-MK2 cells, where a fairly high concentration of active molecules is required for feasible responses in activity. We observe that none of the laboratory mice died after administration of the active extracts at a limited dose during acute toxicity studies. This finding corroborates previous toxicology studies of these plant extracts, which demonstrated no toxicity to laboratory rats [11]. The use of these plants in the treatment of flarial and other infections by the indigenes of Okpambe and other indigenous populations over centuries without serious side effects is proof of their safety. The absence of adverse effects in patients who take the plant concoctions for therapy might be due to *in vivo* detoxification mechanisms, which are absent in *in vitro* studies.

Phytochemical analysis of *S. dulcis* and *C. gabunensis* revealed the presence of secondary metabolites; flavonoids, sterols, phenols, and glycosides in both extracts of the plants,
and alkaloids in the hexane extracts and saponins in the methylene chloride extracts. This suggests that the active principles in the extracts may be from the groups of compounds mentioned above. This is in conformity with previous phytochemical studies, which revealed the presence of these secondary metabolites to be responsible for the medicinal properties of these plants [16, 33].

The isolation of pure compounds from these plant extracts is recommended to determine their full active potential and ascertain the actual phytochemicals responsible for the anti-onchocerca activity of the plant extracts. Such refinement might enhance the activity of the lead compounds against the parasite, and possible reaction mechanisms could be elucidated from any pure lead compound isolated with an attribute of anti-onchocerca activities.

Remarkably, none of the *S. dulcis* and *C. gabunensis* extracts have been previously tested against *Onchocerca* and other *flaria* species. Our data reinforce the existing evidence that these plants have potential antifilarial activity of these plants by traditional health practitioners for the treatment of infectious diseases.

### 5. Conclusions

This study has revealed for the first time the anti-onchocercal activity of extracts of *S. dulcis* and *C. gabunensis*, which could serve as potential novel sources of new drugs against onchocerciasis. It also validates the use of these plants by traditional health practitioners in the local management of filariasis.

### Data Availability

The data used to support the findings of this study are included within the article.

### Conflicts of Interest

The author declares they have no conflicts of interest.

### Authors’ Contributions

Fidelis Cho-Ngwa conceptualized the study. Tiku Edward Tiku and Moses Samje curated the data. Napoleon Mfonku investigated the study. Fidelis Cho-Ngwa and Samje Moses collected the resources. Fidelis Cho-Ngwa supervised the study. Tiku Edward Tiku visualized the study. Tiku Edward Tiku and Samje Moses written the original draft. Tiku Edward Tiku, Samje Moses, and Fidelis Cho-Ngwa edited and reviewed the manuscript.

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