Perforation index assessment of cowpea seeds against cowpea bruchid, *Callosobruchus maculatus* (Fabricius) [Coleoptera: Chrysomelidae], infestation using *Piper guineense*

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**Abstract**

**Background:** Powders and extracts of *Piper guineense* seeds and leaves were assessed for insecticidal activities against *Callosobruchus maculatus* in the laboratory at temperature and relative humidity of 29.6 °C and 75.9%, respectively. Bioactive compounds in *P. guineense* leaves and seeds were also investigated. The powders were tested at rates 1.0, 2.0 and 4.0 g/20 g cowpea seeds while extracts were tested at 1.0, 2.0 and 3.0%.

**Results:** Results of contact toxicity assay of the seed powder caused 100% adult mortality at 96 h post-treatment period whereas leaf powder evoked 90% adult mortality within the same period at concentration of 1.0 g/20 g cowpea seeds. Low adult emergence was observed on cowpea seeds treated with 1 g of seed powder with percentage adult emergence of 0.0% and inhibition rate (IR) of 97.5%. Beetle Perforation Index (BPI) obtained from treated cowpea seeds was significantly different (*P* < 0.05) from BPI of untreated seeds. Extracts of *P. guineense* seed were more toxic than seed powder. *Piper guineense* seed extract caused 87.5% adult mortality of *C. maculatus* while leaf extract caused 70.0% adult mortality within 24 h of infestation at concentration of 1%. Progeny development of *C. maculatus* was completely inhibited in cowpea treated with 2% and 3% leaf and seed extracts of *P. guineense*. β-Pinene was the most abundant active compound in *P. guineense* seed (55.6%) and leaf (48.4%). β-Phellandrene occurred 38.2% in seeds while Ocumene had the least value of 0.2% in seed and 0.5% in leaf.

**Conclusion:** The study showed that *P. guineense* seed powder and extracts were more effective than leaf powder and extract. Utilization of plant products as alternative to synthetic insecticides in protecting cowpea seeds against *C. maculatus* should be encouraged for enhanced food safety and security. *Piper guineense* is used as spice and medicine and interestingly safe for human use.

**Keywords:** Seeds, Insecticidal activities, Leaves, Bioactive compounds, Adult mortality, β-Pinene
Background
There is widespread deficiency in protein and other valuable nutrients in developing countries with bad consequences on the growth and development of children. Cowpea, *Vigna unguiculata* (Walp), seed is palatable and has high content of protein, vitamins and minerals (Nwosu, 2019). The cultivation and consumption of cowpea seeds should be encouraged because its consumption has become necessary with increasing emphasis on plant-based diet (Larochelle, Katungi and Cheng, 2016). For food security, economic and agronomic reasons, cowpea seeds must be stored after harvest and unfortunately in storage, the product is greatly damaged by *Callosobruchus maculatus* (Fab.) (Coleoptera: Bruchidae) insect pest (Casswell, 1981). Prolonged storage of cowpea seeds, particularly at small-scale farming and household levels, is limited due to infestation by *C. maculatus* and this usually culminates in loss in both quality and quantity of the product (Ileke, Odeyemi and Ashamo, 2013). Qualitative and quantitative depletions adversely affect available dietary protein level, market value and specimens for planting during next cropping season, and these are obstacles to achieving food security in many developing countries (IITA, 1995; Rouanet, 1992). The control of *C. maculatus* has relied on the use of synthetic insecticide, which unfortunately has caused problems such as development of resistance, pest resurgence, food poisoning and environmental contamination (Idoko and Adesina, 2012). The ecological and health problems associated with use of chemical insecticides justify the search for alternative methods of controlling *C. maculatus* such as the use of plant materials which are unquestionably safer.

In Nigeria and many other tropical and subtropical countries, there is an array of medicinal plants which could play a fundamental role in pest management (Oparaekwe and Bunmi, 2006). A number of plants used locally for medicinal purposes have demonstrated potential for insect control (Arannilewa, Ekrakene and Akinneye, 2006). These are part of the motivation for our interest in plant materials.

*Piper guineense* (Piperaceae) is a West African species of *Piper* also branded as West African pepper indigenous to Central and Western Africa topical regions and semi-cultivated in Nigeria where the plant leaves are used for flavouring (culinary uses) purposes. Medicinal uses, cosmetic (Dalziel, 1955) and insecticidal potential of *P. guineense* seeds had been reported (Fasakin and Aberjeo, 2002; Idoko & Adesina, 2012). This work therefore is aimed at determining the action of leaf and seed powder products of *P. guineense* in suppressing *C. maculatus* infestation on stored cowpea. Bioactive compounds in *P. guineense* leaves and seeds were also investigated and this is very important in holistic investigations.

Methods
Experimental location
This study was conducted at mean temperature and relative humidity of 29.6 °C and 75.9%, respectively in the Research Laboratory of Animal and Environmental Biology Department, Adekunle Ajayin University, Akungba Akoko (AAUA), Ondo State, Nigeria, and at Insect Chemical Ecology Laboratory, Institute of Bioresources and Sustainable Development, Takyelpat, Imphal, 795001, Manipur, India.

Insect culture
The initial insects used to establish a laboratory colony of *C. maculatus* was obtained from a batch of infested “Oloyin” (Susceptible variety) local cowpea cultivar collected from Ibaka market, Akungba Akoko, Ondo State, Nigeria. The insect culture was by the methods reported by Odeyemi and Daramola (2000) and Ileke et al. (2013). One hundred pairs (10 males to 10 females) of newly emerged adults of *Callosobruchus maculatus* were introduced into the 500 g of cowpea seeds in kilner jar. The kilner jar was then covered with muslin cloth held tightly with rubber band to allow easy flow of air and at the same time prevent the insects from escaping. The insect culture was kept in the laboratory for 1 month (30 days) to allow the insects to oviposit and multiply. The new adults that emerged were subsequently reared on clean uninfested Ife brown variety in the laboratory and serve as the stock culture of the insects used throughout the experiment. The insects were reared under a laboratory condition of 28 ± 2 °C temperature and 75 ± 5% relative humidity.

Plant collection and preparation
Leaves and seeds of *P. guineense* were obtained in fresh form from the herbal stall of Ibaka market, Akungba Akoko, Ondo State, Nigeria, and authenticated at the Plant Science and Biotechnology Department, AAUA, Ondo State, Nigeria. The plant materials were rinsed in clean water to remove dirt and other impurities, cut into smaller pieces, air dried in a well-ventilated laboratory and milled into very fine powder using an electric blender. The resulting powders were then kept in air tight containers and labelled separately inside the refrigerator at 4 °C to maintain their quality before application.

About 150 g of *P. guineense* leaves powder was soaked in an extraction bottle containing absolute methanol for 72 h and the mixture was stirred occasionally with a glass rod. The mixture was stirred occasionally using a glass rod and in order to ensure uniformity in extraction. The extraction process was carried out using a double layer of Whatman No. 1 filter papers. The mixture of the solvent and the extract was separated using a rotary
evaporator at 30 to 40 °C with rotary speed of 3 to 6 rpm for 8 h (Udo, Ekanem and Inyang, 2011). The resulting extracts were then air dried in order to remove traces of the solvent. The same procedure was adopted for the extraction of other plant product (P. guineense seeds). The crude extracts were kept in an amber bottle labelled and preserved in the refrigerator till further use. From this stock solution, 1% concentration was prepared by diluting 0.1 ml of extract in 9.9 ml of solvent; 2% concentration was prepared by diluting 0.2 ml of extract in 9.8 ml of solvent and 3% concentration was prepared by diluting 0.3 ml of extract in 9.7 ml of solvent (Ashamo and Akinnawonu, 2012).

Collection of cowpea seeds
The experimental cowpea seeds (Ife brown variety) were gotten from a newly stocked seeds (without insecticide treatment) in a store at Ibaka market, Akungba Akoko, Ondo State, Nigeria. The seeds were properly sieved, handpicked and disinfested by keeping at −5 °C for 7 days to kill all hidden infestations (if any). All the life stages, particularly the eggs, are very sensitive to cold (Koehler, 2003). The disinfested cowpea seeds were later air dried in the laboratory to prevent mouldiness (Adedire, Obembe, Akinkurolele and Oduleye, 2011) before they were stored in plastic containers with tight lids until ready for use.

Toxicity of P. guineense leaves and seeds powders to C. maculatus
Fine powders of P. guineense leaves and seeds were admixed with cowpea seeds at the rate of 1 g, 2 g and 4 g/20 g of cowpea seeds in 250 ml plastic containers. Ten copulating pairs of C. maculatus (0–3 days old) were introduced into the plastic containers and untreated cowpea seeds replicated four times. The sex of the beetle was determined according to procedures outlined by (Odeyemi & Daramola, 2000). Mortality of adult C. maculatus was evaluated daily for 4 days. Sharp pin was used to probe whether the insect was alive or dead. At 96 h after infestation, all insects were removed from treated and untreated seeds. Percentage adult mortality was corrected using Abbott formula (Abbott, 1925).

\[
\% \text{ Adult Mortality} = \frac{\text{Number of dead adults}}{\text{Number of adults introduced}} \times 100
\]

Oviposition by adult cowpea bruchid on cowpea seeds were recorded before returning the seeds to their respective containers for the emergence of first filial generation. Emerged adult insects were expressed in percentage using standard method as follows:

\[
\% \text{Adult emergence} = \frac{\text{Total number of adult emergence}}{\text{Total number of eggs laid}} \times 100
\]

Progeny inhibition rate (IR) was also calculated using the method described by Tapondjoun, Alder, Bonda and Fontem (2002).

\[
\% \text{ IR} = \frac{C_n - T_n}{C_n} \times 100
\]

where

\[
C_n = \text{number of emerged insects in the control; and}
T_n = \text{number of emerged insects in the treated.}
\]

Reduction in weight of the cowpea seeds was assessed as follows:

\[
\% \text{Weight loss} = \frac{\text{Initial weight} - \text{Final weight}}{\text{Initial weight}} \times 100
\]

Damaged seeds were evaluated by expressing wholesome seeds and bored seeds by adult insect in percentages as follows:

\[
\% \text{Seed damage} = \frac{\text{Number of seeds damage}}{\text{Total number of seeds}} \times 100
\]

Beetle Perforation Index (BPI) was also evaluated for the analysis of damage.

\[
\text{BPI} = \frac{\% \text{treated paddy perforated}}{\% \text{control paddy perforated}} \times 100
\]

BPI value that exceeded 50 will be regarded as enhancement of damaged by cowpea bruchid or negative protectant (Fatope, Mann and Takeda, 1995).

Toxicity of P. guineense leaves and seeds extracts to C. maculatus
The assay follows the same procedure as described for contact toxicity of P. guineense powder above except that 1%, 2% and 3% concentration of each extract of leaves and seeds of the plant material was mixed separately with 20 g of uninfested cowpea seeds in 250 ml plastic containers and this was achieved using a glass rod. Agitation for 10 min then followed immediately to ensure uniform coating. The containers were left open for 30 min to allow solvent traces evaporate. Control experiment was also set up.

Phytochemical screening of P. guineense
Qualitative phytochemical screening were carried out on the methanol extracts of P. guineense seeds and leaves to identify various phytochemical constituents such as alkaloids, tannins, anthraquinones, phlobatannin, cardiac glycosides, saponins and flavonoids present in the plant materials using standard laboratory protocols (Harborne,
1973; Sofowora, 1993; Trease and Evans, 1998; Prashant, Bimlesh, Mandep, Gurpreet and Harleen, 2011).

Gas chromatography-mass spectrometry of *P. guineense*

The GC-MS an analysis of *P. guineense* seed and leaves extracts were performed using a Hewlett-Packard apparatus equipped with an HP-1 fused silica column (30 m × 0.20 mm, film thickness 0.25 μm) and interfaced with a quadrupole detector (Model 5970) operated in electron impact mode. The oven temperature was automated from 70 to 200 °C at 10 °C/min; injector temperature was 220 °C. Helium was used as carrier gas at a flow rate of 0.6 mL/min; the mass spectrometer was operated at 70 eV. Constituent identification was apportioned on the basis of comparison of their retention indices and mass spectra with those given in the Wiley 275 L mass spectral library literature (Adams, 2007).

Statistical analysis

Data were subjected to analysis of variance (ANOVA) and significant treatment means were separated using Tukey’s test. The ANOVA was performed with SPSS 25.0 software (SPSS, 2017).

Results

Mortality of adult *C. maculatus* in cowpea seeds treated with *P. guineense* powders

The effectiveness of the *P. guineense* powders on the survival of *C. maculatus* at different hours after treatment is presented in Table 1. There was a significant difference (*P < 0.05*) in insect adult mortality amongst the treatments. Mortality of insect increased gradually with time of contact with *P. guineense* seed powder causing 100% mortality within 96 h of exposure, whereas the leaf powder evoked 90% mortality within the same period at concentration 1 g/20 g of cowpea seeds. Mortality of *C. maculatus* varied with plant parts, dosage and period of exposure. The highest mortality of 95% was obtained at 3 g/20 g of cowpea seeds with the seed powder within 24 h of exposure, while the leaf powder caused 82.5% mortality of adult insect within 24 h of exposure.

Effect of *P. guineense* powders on *C. maculatus* emergence

Plant powder reduced fecundity of adult *C. maculatus* (Table 2). The amount of eggs laid on treated cowpea seeds by *C. maculatus* was significantly (*P < 0.05*) lower than amount of oviposition on untreated cowpea seeds. There was no significant difference (*P > 0.05*) in the mean number of oviposition cowpea seeds treated with *P. guineense* seed and leaf powders, but were significantly different when compared to untreated seed. The % adult emergence in the control experiment (64%) was significantly higher (*P < 0.05*) compared to cowpea seeds treated *P. guineense* leaf powder (20%) at rate 1 g/20 g of cowpea seeds. However, the lowest number of adult emergence was observed on cowpea seeds treated with seed powder at rate 1 g/20 g of cowpea seeds with percentage adult emergence of 0.0% and inhibition rate (IR) of 100%. Similarly, *P. guineense* leaf powder completely inhibited adult emergence at rate 3 g/20 g of cowpea seeds.

Beetle perforation index caused by *C. maculatus* in cowpea seeds treated with *P. guineense* powders

*Piper guineense* seed powder completely protected cowpea seeds from being damaged by *C. maculatus* concentrations 2 g and 3 g/20 g (Table 3). There was neither seed damage nor weight loss recorded in the treated cowpea seeds. Meanwhile, leaf powder significantly suppressed infestation as manifested in percentage seed damage and weight loss compared to untreated cowpea seeds. Beetle Perforation Index (BPI) was zero in cowpea seed treated with seed powder at rate 2 g and 3 g/20 g of cowpea seeds, leaf powder recorded 10% and 2.5% at rate 1 g and 2 g/20 g of cowpea seeds, while untreated seed recorded > 50.00% perforation index. However, the BPI obtained from treated cowpea seeds was

| *P. guineense* powders | Conc. (g) | % mortality ± SE mean |
|------------------------|----------|-----------------------|
|                        | 24 h     | 48 h                  | 72 h                  | 96 h                  |
| Leaf 1.0               | 50.00 ± 3.25<sup>b</sup> | 77.50 ± 3.25<sup>b</sup> | 82.50 ± 3.25<sup>b</sup> | 90.00 ± 3.04<sup>b</sup> |
| Seeds 1.0              | 70.00 ± 3.25<sup>c</sup> | 82.50 ± 3.75<sup>b</sup> | 95.00 ± 3.50<sup>c</sup> | 100.00 ± 0.00<sup>b</sup> |
| Leaf 2.0               | 75.00 ± 3.50<sup>d</sup> | 80.00 ± 3.04<sup>d</sup> | 90.00 ± 2.04<sup>c</sup> | 100.00 ± 0.00<sup>b</sup> |
| Seeds 2.0              | 87.50 ± 3.25<sup>c</sup> | 95.00 ± 3.50<sup>c</sup> | 100.00 ± 0.00<sup>d</sup> | 100.00 ± 0.00<sup>d</sup> |
| Leaf 3.0               | 82.50 ± 3.75<sup>d</sup> | 90.00 ± 3.04<sup>c</sup> | 100.00 ± 0.00<sup>d</sup> | 100.00 ± 0.00<sup>d</sup> |
| Seeds 3.0              | 95.00 ± 3.50<sup>d</sup> | 100.00 ± 0.00<sup>d</sup> | 100.00 ± 0.00<sup>d</sup> | 100.00 ± 0.00<sup>d</sup> |
| Untreated 0.0          | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> |

Each value is a mean ± standard error of four replicates. Means within the same column followed by the same letter are not significantly different at *P > 0.05* using Tukey’s test.
significantly different from BPI of the untreated. In the untreated cowpea seeds, 42.50% damage occurred as revealed by adult emergent holes of the bruchid. As a result of the feeding activity of *C. maculatus* larvae on the cowpea seeds, the weight of the untreated cowpea seeds was significantly (*P* < 0.05) reduced compared with the treated seeds.

**Mortality of adult *C. maculatus* in cowpea seeds treated with *P. guineense* extracts**

*Piper guineense* seed extract caused 100% adult mortality of *C. maculatus* while leaves extract caused 92.5% adult mortality within 24 h of infestation at concentration 1%/20 g of cowpea seeds. The adult mortality observed in the study with cowpea seeds treated with leaves and seeds extracts increased with the duration of exposure (Table 4) and the mortality recorded was significantly higher compared to solvent treated (+ve control) and untreated (−ve control) seeds. The untreated seeds exerted no adult mortality effects on the adult beetles over the duration of exposure.

**Effect of *P. guineense* extracts on *C. maculatus* emergence**

The plant extract significantly reduced the number of eggs laid by *C. maculatus* compared with untreated cowpea seeds. Cowpea bruchid laid the lowest mean number of eggs (0.0 eggs) on cowpea treated with seed extract followed by 1.00 eggs laid on cowpea treated with leaves extract at concentration 3%/20 g of cowpea seeds (Table 5). *Callosobruchus maculatus* adult emergence at concentrations 2% and 3%/20 g of cowpea seeds was completely inhibited in cowpea treated with both leaves and seed extracts of *P. guineense* with inhibition rate (IR) of 100% compared to control that recorded 64% adult emergence. Treating cowpea seeds with methanol did not prevent progeny emergence from seeds.

| Table 2 Progeny development of *C. maculatus* in cowpea seeds protected with *P. guineense* powders |
| --- |
| **P. guineense powders** | Conc. (g) | Mean number of eggs laid | % adult emergence | %IR |
| Leaf | 1 | 20.00 ± 2.04<sup>ab</sup> | 20.00 ± 2.04<sup>a</sup> | 90.00 ± 3.04<sup>a</sup> |
| Seed | 10.00 ± 1.04<sup>bc</sup> | 10.00 ± 1.04<sup>bc</sup> | 97.50 ± 2.50<sup>b</sup> |
| Leaf | 12.50 ± 1.75<sup>ab</sup> | 8.00 ± 0.04<sup>b</sup> | 97.50 ± 2.50<sup>b</sup> |
| Seed | 7.50 ± 0.25<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 100.00 ± 0.00<sup>a</sup> |
| Leaf | 3 | 7.50 ± 0.25<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 100.00 ± 0.00<sup>a</sup> |
| Seed | 5.00 ± 0.04<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 100.00 ± 0.00<sup>a</sup> |
| Untreated | 0.0 | 62.50 ± 3.75<sup>a</sup> | 64.00 ± 3.22<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> |

Each value is a mean ± standard error of four replicates. Means within the same column followed by the same letter (s) are not significantly different at *P* > 0.05 using Tukey’s test.

| Table 3 Perforation Index caused by *C. maculatus* in cowpea seeds treated with *P. guineense* powders |
| --- |
| **P. guineense powders** | Conc. (g) | Total no of seeds | Number of damaged seeds | % seed damaged | % weight loss | Beetle Perforation Index (BPI) |
| Leaf | 1.0 | 95.25 | 4.00 | 4.50 ± 0.07<sup>ab</sup> | 10.00 ± 1.04<sup>a</sup> | 10.00 ± 1.04<sup>a</sup> |
| Seed | 94.75 | 1.00 | 1.05 ± 0.02<sup>ab</sup> | 5.00 ± 0.02<sup>a</sup> | 2.50 ± 0.02<sup>a</sup> |
| Leaf | 2.0 | 94.50 | 1.00 | 1.05 ± 0.01<sup>ab</sup> | 5.00 ± 0.02<sup>a</sup> | 2.50 ± 0.02<sup>a</sup> |
| Seed | 95.25 | 0.00 | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> |
| Leaf | 3.0 | 94.00 | 0.00 | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> |
| Seed | 95.50 | 0.00 | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> | 0.00 ± 0.00<sup>a</sup> |
| Untreated | 0.0 | 95.25 | 40.00 | 42.50 ± 3.75<sup>a</sup> | 75.00 ± 3.20<sup>a</sup> | >50.00 ± 0.00 |

Each value is a mean ± standard error of four replicates. Means within the same column followed by the same letter are not significantly different at *P* > 0.05 using Tukey’s test.

* - Perforation Index
untreated cowpea seeds was significantly ($P < 0.05$) reduced compared with the treated seeds. The BPI values obtained on seeds treated with 1%, 2% and 3% methanol were significantly higher than 50% compared with seeds treated with plant extracts.

**Phytochemical screening of *Piper guineense***

The results of the qualitative phytochemical constituents of both extracts, as shown in Table 7, revealed the presence of alkaloids, saponins, tannins, flavonoids and cardiac glycosides, while phlobatannins and anthraquinones were not present in both leaves and seeds of the plants.

**Bioactive compounds *P. guineense* leaves and seeds**
The major bioactive compounds identified by GC-MS analysis from the leaves and seeds of *P. guineense* are shown in Table 8. The result shows that both seeds and leaves contain the same bioactive compounds at varying percentages. β-Pinene was the most abundant 55.6% in seeds and 48.4% in leaves of *P. guineense*. This was followed by β-Phellandrene which occurred 38.2% in seeds while the least was Ocimene with a value of 0.2% in seeds and 0.5% in leaves.

**Discussion**

Plants are rich source of bioactive chemical compounds with insecticidal properties (Rajkumar and Jebanesan, 2004). The activity of plant products is ascribed to the complex mixture of active compounds (Sukhthankar, Kumar, Godinho and Kumar, 2014). The potential of powdered and methanol extract of *P. guineense* leaves and seeds in causing acute toxicity, preventing oviposition and perforation to cowpea seeds by *C. maculatus* has been demonstrated in this study and results obtained conformed with the conclusions of Su (1977), Ivbijaro and Agbaje (1986) and Ogunwolu, Igo and Longs.
(1988) who reported insecticidal activity of *P. guineense* extract and powdered products against *Sitophilus oryzae* and *Callosobruchus maculatus*, respectively.

The effectiveness of *P. guineense* on mortality of cowpea bruchid in this work is due to contact toxicity effect as a result of volatile constituent which may lead to respiratory impairment, which affects their metabolism and consequently other systems of the insect body (Islam et al., 2009; Ofuya and Dawodu, 2002). Ogunwolu et al. (1988) viewed that *P. guineense* powder can cause physical abrasion of the insect cuticle which can lead to loss of body fluid or blockage of spiracles resulting into suffocation of the insect. The significant insect mortality caused by the application of seeds powder and extracts may be ascribed to the presence of highly spicy alkaloidal secondary metabolites (Udo et al., 2011). This prevented locomotion, physical contact of adult beetles with seeds and triggered asphyxia, starvation of adult insect or unknown physiological changes activation (Adesina, Jose, Rajashekar and Afolabi, 2015).

The reduced oviposition and completely inhibited progeny development clearly indicated that *P. guineense* mechanism of action is by oviposition deterrence and toxicity to eggs (ovicidal). The suppression of oviposition by the insects in treated containers compared to untreated may be due to locomotion impediment; hence, the beetle was unable to move freely thereby affecting mating activities and sexual communication (Adesina, 2013; Ileke, 2014; Odeyemi & Daramola, 2000) resulting in few number of eggs observed in this study. Zabri, Kabran, Kodjo and Trabi (2009) opined that the ability of pesticidal plant products to reduce the egg laying capability by female beetles may be ascribed to the occurrence of flavonoids (Ileke, 2014). The reduction in oviposition in the extract-treated seeds compared to powder may also be caused by an

### Table 6 Perforation Index caused by *C. maculatus* in cowpea seeds treated with *P. guineense* powders

| *P. guineense* extracts | Conc. (% | Total no of seeds | Number of damaged seeds | % seed damaged | % weight loss | *Beetle Perforation Index* (BPI) |
|-------------------------|---------|--------------------|--------------------------|---------------|--------------|--------------------------------|
| *P. guineense* leaf     | 1       | 93.50              | 1.00                     | 2.50 ± 0.02ab | 5.00 ± 0.04a | 5.50 ± 0.05c                   |
| *P. guineense* seeds    | 95.25   | 0.00               | 0.00                     | 0.00 ± 0.00a  | 0.00 ± 0.00a | 0.00 ± 0.00a                   |
| Methanol (+ve control)  | 92.00   | 0.00               | 0.00                     | 32.50 ± 2.25cd| 40.00 ± 2.04c| 75.00 ± 3.04c                   |
| *P. guineense* leaf     | 2       | 94.00              | 0.00                     | 0.00 ± 0.00a  | 0.00 ± 0.00a | 0.00 ± 0.00a                   |
| *P. guineense* seeds    | 94.02   | 0.00               | 0.00                     | 0.00 ± 0.00a  | 0.00 ± 0.00a | 0.00 ± 0.00a                   |
| Methanol (+ve control)  | 96.00   | 0.00               | 0.00                     | 27.50 ± 2.50c | 37.50 ± 2.50c| 65.00 ± 3.04c                   |
| *P. guineense* leaf     | 3       | 95.00              | 0.00                     | 0.00 ± 0.00a  | 0.00 ± 0.00a | 0.00 ± 0.00a                   |
| *P. guineense* seeds    | 94.50   | 0.00               | 0.00                     | 0.00 ± 0.00a  | 0.00 ± 0.00a | 0.00 ± 0.00a                   |
| Methanol (+ve control)  | 93.00   | 0.00               | 0.00                     | 25.00 ± 2.20c | 35.00 ± 2.20c| 60.00 ± 3.04c                   |
| Untreated (--ve control)| 0.0     | 95.25              | 40.00                    | 42.50 ± 3.75d | 52.50 ± 3.20d| >50.00 ± 0.00                   |

Each value is a mean ± standard error of four replicates. Means within the same column followed by the same letter are not significantly different at P > 0.05 using Tukey’s test.

* - Perforation Index

### Table 7 Phytochemicals in different extracts of *P. guineense* leaf and seed

| Phytochemicals | Methanol extract of *P. guineense* leaf | Aqueous extract of *P. guineense* leaf | Methanol extract of *P. guineense* seed | Aqueous extract of *P. guineense* seed |
|----------------|----------------------------------------|---------------------------------------|----------------------------------------|---------------------------------------|
| Alkaloids      | +                                      | +                                     | +                                      | +                                     |
| Saponins       | +                                      | +                                     | +                                      | +                                     |
| Tannins        | +                                      | +                                     | +                                      | +                                     |
| Phlobatannins  | –                                      | –                                     | –                                      | –                                     |
| Anthraquinones | –                                      | –                                     | –                                      | –                                     |
| Flavonoids     | +                                      | +                                     | +                                      | +                                     |
| Cardiac glycosides | +                | +                                     | +                                      | +                                     |

– negative, + positive
Table 8 Major bioactive chemical composition of *P. guineense* leaves and seeds extracts

| Bioactive molecules | Retention time | Leaves (%) | Seeds (%) |
|---------------------|----------------|------------|-----------|
| β-Phellandrene      | 10.12          | 3.4        | 38.2      |
| a-Phellandrene      | 10.12          | 4.5        | 7.2       |
| Limonene            | 10.20          | 15.7       | 10.3      |
| Ocimene             | 10.45          | 0.5        | 0.2       |
| Linalol             | 10.90          | 1.5        | 4.6       |
| a-Pinene            | 12.20          | 6.3        | 10.3      |
| Pyridine            | 12.45          | 1.4        | 2.3       |
| β-Pinene            | 12.70          | 48.4       | 55.6      |
| β-Myrcene           | 13.26          | 1.4        | 2.5       |
| a-Cubebeine         | 13.52          | 3.0        | 1.0       |
| α-Copaene           | 13.78          | 0.7        | 1.2       |
| α-Caryophellene     | 14.19          | 4.1        | 17.6      |
| α-Curcumene         | 14.80          | 1.2        | 1.0       |
| (E, Z) α-Foreseen   | 15.21          | 3.9        | 1.9       |
| cis-α-Bisabolene    | 15.00          | 1.0        | 0.6       |
| Eugenol             | 19.89          | 1.3        | 3.7       |
| Piperanol           | 20.39          | 3.2        | 7.9       |

The significant inhibition of the progeny development of *C. maculatus* by the seed powder and extract of *P. guineense* could also probably be attributed to the presence of appreciable vapour pressure from which poison toxic amounts could be ingested picked by insects through the vapour phase (Lale, 2002). The pungent vapour of *P. guineense* might have diffused into seeds and ingested by the larval stages (moulting) of *C. maculatus* developing within the seed thus leading to reduced or complete inhibition of progeny emergence due to impaired physiological and biochemical process associated with post-embryonic development (Ketoh, Koumaglo, Glitho and Huignard, 2006; Udo et al., 2011).

Untreated cowpea seeds suffer great damage due to *C. maculatus* infestation, whereas negligible or no damage/weight loss were recorded in cowpea seeds that were treated with the *P. guineense* leaves powders and extract. Beetle Perforation Index (BPI) value lower than 50 is an index of positive protectant effect while BPI greater than 50 is an index of negative protectability. The reduction of damage observed in this study is the consequence of the higher adult mortality, antifeedant, oviposition deterrence, ovicidal, larvical and reproduction inhibitory properties of *P. guineense* (Isman, 2006; Lale and Abdulrahman, 1999; Manikanta and Dokuparthi, 2014). The efficiency of seed extract obtained was in consonance with the findings of Eziah, Buxton and Owusu (2013) who reported that methanol extracts of *Zanthoxylum zanthoxyloides* and *Securidaca longepedunculata* roots significantly reduced the damage caused by *Prostephanus truncatus* and *Tribolium castaneum* on maize grains.

Insecticidal activity of plant materials depends on the abundance of active compounds of the plant material. The major bioactive constituents isolated from the methanol leaves and seeds extracts of *P. guineense* were similar to those reported in *P. guineense* obtained from Cameroun, South Eastern and South Western Nigeria (Oben, McConchi, Phan-Thien and Ntonifor, 2015; Ojinnaka, Ubbor, Okudu and Uga, 2016; Owolabi, Lawal, Ogunwande, Hauser and Setzer, 2013; Oyediji, Adeniyi, Ajayi and König, 2005; Tchoumgoungnang et al., 2009); however, there was a slightly difference in the concentrations of the major bioactive constituents found. The observed compositional variability in the bioactive chemical composition might be due to several factors such as environmental and climatic, soil fertility variations, genetic makeup of the plant, phenological state of the plant, different chemotypes and solvents used for extraction (Isman and Machial, 2006; Perry et al., 1999). The screening of this plant species revealed the presence of tannins, saponins, alkaloids, flavonoids and cardiac glycosides. It was reported that *P. guineense* contains bioactive ingredients such as neurotoxic piperamides and lignans, alpha-pinene, limonene, Linalool, piperine and chavicine, which are insecticidal including piperidine and alkaloids as the major active components in *P. guineense* seeds (Golob, Mwumbola and Mbhango Ngulube, 1999; Lale, 1995; Scott et al., 2004; Scott, Gagnon, Lesage, Philoge and Arnason, 2005). The plant phytochemicals exerted their action on the physiological, biochemical and enzymatic process of the insects. Many studies have identified β-Phellandrene, eugenol, limonene, linalool, α-pinene and β-pinene as components of plant essential oils and also tested fumigant and extract film on the seeds which makes it becomes unsuitable for oviposition (Adesina and Ofuya, 2015).

The experimental plant products significantly reduced adult emergence or progeny development in treated cowpea seeds. The reduction in adult emergence could be due to ovicidal or larvicidal effects of the plant products. The results from the study shows that few eggs were oviposited on cowpea seed treated with powder and extract of *P. guineense* seeds but with no adult emergence. Partial blastokinesis and abnormal breakage of extra embryonic membranes in the embryo can result into reduction in progeny development or emergence of teneral adult insect (Enslee and Riddiford, 1977). The ability of the evaluated plant materials to significantly suppress or inhibit adult emergence confirms the insecticidal activity of the plant species in reducing oviposition and adult emergence (Abdullahi and Muhammad, 2004).
contract toxicities of phytochemicals from plant (Beyrouthy et al., 2011; Demirci, Özek and Baser, 2000; Nenaah, 2014; Phillips and Appel, 2010; Phillips, Appel and Sims, 2010; Yeom et al., 2015). However, there are scarce reports on their pesticidal potential because they are not commercialised (Szolyga, Gnilka, Szczerpik and Szuny, 2014).

Conclusion

This study has further demonstrated that *P. guineense* leaf and seed possess phytochemicals that confer on it significant insecticidal value as a potential stored seeds protectant, having the capacity to evoke high mortality within 24 h and greatly suppressed oviposition, progeny development and seed damage. Its adoption as suitable alternative to synthetic insecticides as seed protectant should be encouraged amongst resource-poor farmers as a means of ensuring a steady supply of quality food, since it has traditionally been used as a spice and medicine and has been proven safe for humans.

Abbreviations

ANOVA: Analysis of variance; IR: Progeny inhibition; BPI: Beetle Perforation Index; GC-MS: Gas chromatography-mass spectrometry; IITA: International Institute for Tropical Agriculture; +: Positive; -- Negative

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Authors’ contributions

This research work was carried out in collaboration amongst all authors. KDI conceived and designed the study and contributed to the data analysis and manuscript reviewing; KDI, AO and LCN collected the data on insect bioassay; JMA conducted the GC-MS analysis and reference search and contributed to the manuscript draft. All authors read and approved the final manuscript.

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