

Original Research Article

The Climbing Performance, Neuromuscular transmitter (AChE) Activity, Reproductive Performance and Survival of *Drosophila melanogaster* Fed Diet with *Mangifera indica* Cold Aqueous Leaf Extract

Abstract

Objective: To screen the toxic effect of *Mangifera indica* aqueous leaf extract in *Drosophila melanogaster*.

Materials and Methods: Phytochemical screening was carried out. 20 Adult Flies were exposed to 7.5 mg, 15 mg, 30 mg, 45 mg and 100 mg /10 g diet for acute toxicity (168hrs) while 50 flies were exposed to 2.5 mg 5 mg and 10 mg/10 g diet for sub chronic (28 days). All concentrations were prepared in 1000ul of distilled water and replicated three (3) times. Diet+1000µl of Distilled water served as control. Fecundity/developmental toxicity, Climbing and AChE activities were carried out by exposing Flies to the sub-chronic concentrations for 5 days.

Results: Phytochemical screening revealed the presence of Alkaloids, flavonoids, tannins, saponins and terpenoids. The acute toxicity test showed 100% mortality at 100 mg/ 10 g diet and 168hrs LC₅₀ was 72.4 mg/10 g diet. The sub chronic toxicity test showed decrease in Flies survival along concentration with a least survival at 10 mg/10 g diet. There was a slight reduction and elevation in the Climbing and AChE activities respectively but not statistically significant ($p>0.05$) compared to control. At 5 mg and 10 mg/10 g diet there was a delay in the development with few enclosed Flies.

Conclusion: from the Results, it can be concluded that *Mangifera indica* aqueous leaf extract may be toxic at high dose from 72.4 mg/10g diet and might have an adverse effect on the development and Survival of flies at sub chronic concentration as low as 2.5mg/10g diet.

Chapter One

1.0 Introduction

From time immemorial, nature has been of great help to humans and animals alike, particularly plants that have been a source of new drugs with curative properties to treat various diseases. Plants are used in traditional medicine to treat different infectious and noninfectious diseases [1, 2]. These treatments include body-washes, massages, ingestions, etc. [3]. It has been documented that nearly 80% of the world's population relies on traditional medicines for primary health care, most of which involve the use of plant extracts [4]. Little or no detailed information on the toxicological profile of these medicinal plants has been documented; some plant extracts can be hepato-toxic, neurotoxic, hemotoxic and damaging to the reproductive cells even at lower concentration; despite the medicinal properties of these plant extracts against infections and diseases. Reproductive fitness is a broad, ecological measure of health [5]. Many herbs can negatively affect male and female fertility as well health during pregnancy, [6]. Some medicine can alter ovulation and endometrial or uterine receptivity to a pregnancy, [7]. Medical plant can however be poisonous if wrong plant part or wrong concentrations are used [8]. Now a days, toxicity and safety of medicinal plants are the most concerning topics, as use of plant product has become popular worldwide [9]. Toxicity can be Acute, sub-chronic and chronic depends on the period and concentration of exposure. It has been shown that the toxicity of a given plant depends on various factors, including the strength of secondary metabolites, the quantity consumed, the time of exposure, different parts of the plant (root, oil, leaves, stem bark and seeds), individual body chemistry, climate and soil, and genetic differences within the species [10] Indeed, of about 1,500,000 plants investigated, most of them contain toxic substances [11]

Mangifera indica is a large evergreen and a heavy, dome-shaped crown tree. It is in the family Anacardiaceae [12]. It is distributed all over the tropical regions of the world [12]. It has been useful as horticultural and medicinal plant in this tropical region of the world. Fruits contain protein, fat, carbohydrate, minerals, vitamins A, B and C and amino acids. The fruits also yield a resin containing mangiferene, mangiferic acid, resinol and maniferol and others [13, 14]. The leaves have been reported to contain saponins, glycosides, unsaturated sterols, polyphenols, euxanthin acid, mangiferine and tannins etc. Burns, scalds, sores, cough and diarrhoea are treated with the ashes of the leaves of *Mangifera indica* in South America and other parts of the world [13, 15]. The *in vivo* protective antioxidant activity of *Mangifera indica* in *Drosophila melanogaster* has been reported [12]. Antiseptic

activity leaf extracts in the treatment of infections, wounds scalds, burns, sores, and abscesses in animals and humans has been documented in a number of ethnobotanical surveys [16].

To study the toxicity of medicinal plants, the *in-vitro* and *in-vivo* models are available. Regardless of the type of extract, the parts of plant used, the concentration of the extract, the mode of administration, and the organism under consideration the lethal dose/concentration 50 (LD/LC50); the dose/concentration that can kill 50% of a tested population is used to establish the toxicity of the plant extract [17].

The use of an *in vivo* approach facilitates the crucial understanding of how chemicals affect complex metabolic targets and pathology at the cell and organ level. Mice, rats, rabbits, and other laboratory mammals have been used extensively to study chemical toxicity. However, *in vivo* assays with those species are often expensive, use large amounts of toxic test chemicals, and are difficult to use in a moderate-throughput to high-throughput manner. Those drawbacks have led toxicologists to develop alternative animal models for chemical testing. Many alternative test organisms share biological processes with rodents and other mammals, including humans. Three test platforms of note that can be adapted to high-throughput screening rely on insect, nematode, and zebrafish models [18]

Drosophila melanogaster, known colloquially as the fruit fly, remains one of the most commonly used model organisms for biomedical science [12]. Due to the success achieved in the use of *D. melanogaster* in experimental studies, it has met the standard of the European Centre for the Validation of Alternative Methods (ECVAM): Reduction, Refinement and Replacement (3Rs) of laboratory animal usage [19]. For more than one hundred years, the low cost, rapid generation time, and excellent genetic tools have made the fly indispensable for basic research [12]. *Drosophila* is also a good model for studying toxicological effect on multiple generations owing to their short life cycle, valuable for the analysis of molecular mechanisms underlying genetic phenomena, behavior and development. Approximately 65% of human disease-causing genes are believed to have a functional homolog in flies [20] and a significant fraction of these homolog are expressed in *Drosophila* tissues that perform the function of the equivalent human tissue [21]. In addition to similarities in basic cellular structure and function, humans and *Drosophila* share pathways for intercellular signaling [22], learning and behavior [23] etc. Fruit flies have the potential to be used for chemical-toxicity screens [24, 25].

The use of *Mangifera indica* extract in the treatment of diseases in the western part of Nigeria is on the increase and the dearth of information on the acute and sub-chronic toxicity of this plant is becoming a course of concern. Therefore, it is necessary to evaluate the toxicity of *Mangifera indica* cold aqueous leaf extract in *Drosophila melanogaster*.

Chapter two

2.0 Plant Collection

The leaf of *Mangifera indica* was collected from University Senior Staff Quarters, Jos, Plateau State, Nigeria. It was identified by a Plant taxonomist in plant science Department University of Jos. It was air dried using room temperature for 7 days, pulverized using mortar and pestle and extracted using distilled water 1:10 w/v as described in our previous work [12]

2.1 Animal Model

D. melanogaster Harwich strain was obtained from the Africa Centre of Excellence in Phytomedicine Research and Development. Species Stock was maintained at constant temperature and humidity (23 ± 1 °C; 60% relative humidity, respectively) under 12 h dark/light cycles. We used standard *Drosophila* medium composed of cornmeal medium (1% w/v), brewer's yeast (2% w/v), agar and nipagin (0.08% w/v). The maximum average lifespan of flies used here (*D. melanogaster*, Harwich strain) is about 50–58 days, and about 50% of the flies generally die within 41–45 days. However, this can vary depending on a variety of factors such as diet composition and temperature (26).

2.2 Phytochemical Screening

Phytochemical examinations of *Mangifera indica* cold aqueous leaf extract was carried out to determine the presence of metabolites such as alkaloids, flavonoids, phenols, saponins, terpenoids, tannins and glycosides according to standard methods [27]. Any visible change of color or precipitate formation was used as an indicative for the presence (+) or absence (–) of individual metabolites

2.3 Experimental Design

The Leaf extract of *Mangifera indica* concentration, 7.5 mg, 15 mg 30 mg, 45 mg and 100 mg were prepared in 1000ul of distilled water and 9g of diet given each concentration per 10g diet (7.5 mg/10 g diet, 15 mg/10 g diet, 30 mg/10 g diet 45 mg/10 g diet and 100 mg/10 g diet). Diet with 1000ul of Distilled water served as control. 20 adult Flies were exposed to these concentrations for 7 days and 168 hrs LC₅₀ Acute concentrations [were](#) evaluated. For sub-chronic (survival) evaluation, 2.5 mg, 5 mg and 10 mg concentration of *Mangifera indica* cold aqueous leaf extract were prepared in 200 ul of distilled water and 9.8 g diet making a total of 10g (200 ul of D.water + 9.8g diet) and were exposed to flies for 28 days. Five (5) Days-Short term exposure of *D. melanogaster* to the sub-chronic concentration (2.5 mg/10 g diet, 5 mg/10 g diet and 10 mg/10 g diet) was carried out to evaluate the Climbing performance, Acetylcholinesterase activity and fecundity of the exposed flies.

2.3.1 Determination of 168 hrs LC50

This study was carried out following the method described by Sayanti & Sumedha [28], with slight modification. 20 flies of age rang 1- 3 days were anesthetized, counted and exposed to graded concentration of *Mangifera indica* cold aqueous leaf extract. The following Concentrations, 7.5 mg/10 g diet, 15 mg/10 g diet, 30 mg/10 g diet 45 mg/10 g diet and 100 mg/10 g diet for the first phase. The second phase concentration ranging from 500 mg, 1000 mg and 2000 mg/10 g diet. Each phase is for 168 hrs (7 days). Mortality reading was scored every 24 hrs interval for 7 days. The mortality rate was subjected to Probit analysis of Graphpad prism 7.04 for LC₅₀ determination.

2.3.2 Survival of *Drosophila melanogaster* Fed *Mangifera indica* Aqueous Leaf Extract

In this experiment, 50 flies of both genders (1-3 days old) were exposed to 2.5mg/10 g diet , 5mg/10g diet and 10mg/10g diet of *Mangifera indica* aqueous leaf extract in five replicates for 28 days as described by [29]. The number of live and dead flies was scored daily till the end of the experiment and the survival rate was expressed as percentage of live flies. Short term exposure of subchronic concentration was also set up for 5 days in triplicates, 50 adult flies each and were used for climbing assay activity, fecundity assay and Homogenized in 0.1m PBS pH 7.4, centrifuged using cold centrifuge. The supernatant was used for determining Acetyl holiness erase activity of the exposed flies.

2.3.3 Negative Geotaxis (Behavioral Assay)

The locomotor (Climbing) performance of *Mangifera indica* aqueous leaf extract exposed and control flies would be investigated using the negative geotaxis assay [29]. Briefly, ten (10) *Mangifera indica* aqueous leaf and bark stem extract exposed and control flies would be immobilized under mild ice anesthesia and placed separately in labeled vertical glass columns (length, 15 cm; diameter, 1.5 cm). After the recovery period (about 20 min), the flies would be gently tapped to the bottom of the column. Following 6 s, the number of flies that climb up to the 6 cm mark of the column, as well as those that remain below this mark would be recorded. Data would be expressed as the percentage of flies that escape beyond the 6 cm mark in 6 s. The score of each group is an average of three trials for each group of treated and controlled flies.

2.3.4 Fecundity Assay

Paragraph Virgin fruit flies (Five females) with five males were collected as described by Berg laboratory. They were allowed to feed on *Mangifera indica* cold aqueous leaf extract containing diet for 5 days. Males and **female** were treated separately (separate vials) for this exposure period of 5 days. They were combined and allowed to mate on a normal diet without an extract for 24 hours. After 24 hours the adult flies were transferred off diet and sacrificed. The **Vials** were maintained at constant temperature and humidity (23±1 °C; 60% relative humidity, respectively) under 12 h dark/light cycles for the development of egg laid for 12 days. The number of emerged adults was recorded.

2.3.5 Acetylcholinesterase (AChE) activite

Eliminate comma

The 5 days *Mangifera indica* cold aqueous leaf extract-treated flies, were **anesthetize** on ice, homogenized with 100mM phosphate buffer saline pH 7.4 and centrifuged using 4°C Cold centrifuge for 10 min at 3000 rpm. The supernatant was collected and used for the determination of AChE activity following the method described by Ellman et al. [30] with slight **Modification**. To the reaction mixture containing 285 ul of distilled water, 180 ul of 100 mM potassium phosphate buffer (pH 7.4), 60 ul of 10 mM DTNB, and 15 ul of sample, 60 ul of 8 mM acetylthiocholine was added. The change in absorbance was monitored at 412 nm for 2min at 15 s intervals, using a UV-VIS Spectrophotometer (Jenway). The data were calculated against blank, **and** sample blank and the results were corrected by the protein content. The enzyme activity was expressed as mmole/min/mg of protein.

Chapter Three

3.0 Results

3.1 Qualitative Phytochemical Screening

Phytochemical examinations of *Mangifera indica* cold aqueous leaf extract revealed the presence of alkaloids, flavonoids, tannins, saponins and terpenoids while glycoside were absent. The result is presented in table

1.

Test	Results
Alkaloids	++
Flavonoids	+++
Phenols	++
Tannins	++
Saponins	+
Terpenoids	+
Glycosides	-

Absent (-), slightly present (+), moderately present (++) and highly present (+++)

Table1: [Qualitative](#) Phytochemical Screening Result of *Mangifera indica* cold aqueous extract

[Qualitative or quantitative? Or Qualitative and quantitative?](#)

3.2 168 hrs LC₅₀

The mortality rate of *Mangifera indica* cold aqueous leaf extract was 4% and 95% at 0 mg/10 g diet (control) and 100 mg/10 g diet respectively. The 168 hrs LC₅₀ was evaluated to be 72.4 mg/10 g diet as shown in figure 1. **Just for standardization , use capital letter or not.**

3.3 28 Days (Sub-Chronic) Survival

The Survival proportion of flies treated with Sub-chronic concentrations, 2.5 mg/10 g diet, 5 mg/10 g diet and 10 mg/10 g diet after 28 days ranged from 2-30%. The highest survival proportion was recorded in the controlled flies while the least was recorded in 10 mg/ 10 g diet. At 2.5 mg/ 10 g diet and 5 mg/10 g diet, the survival proportion was recorded to be 27 % and 7 % respectively. There was significant difference ($p < 0.05$) comparing the survival curve of the treated flies to the control survival curve. Log-rank (Mantel-Cox) test p Value < 0.0001 . The result is presented in figure 2.

3.4 Climbing Activity

The Negative geotaxis of *Mangifera indica* cold aqueous Leaf extract-treated flies ranged from 62-71 %. The highest climbing activity was recorded in the controlled group while the least was recorded in 10 mg/10 g diet. The climbing performance of flies treated with 2.5 mg/10 g diet and 5 mg/10 g diet was recorded to be 70% and 68% respectively. There was no significant difference ($p > 0.05$) comparing the climbing activity of the treated flies to control flies. The result is presented in figure 3

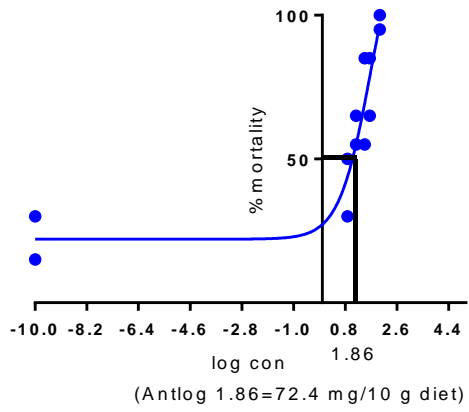


Fig 1: LC₅₀ of *Mangifera indica* Cold aqueous leaf extract in Flies

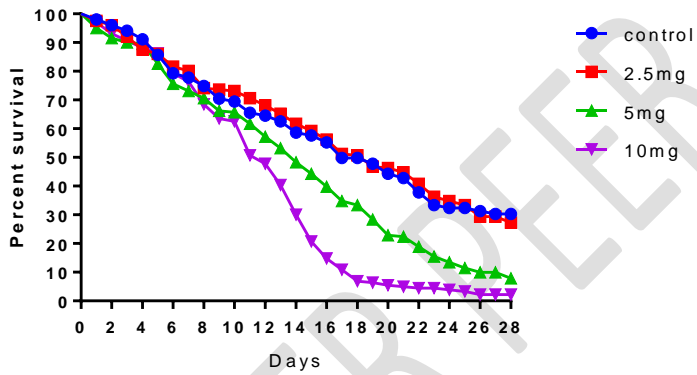


Fig 2: 28 Days (Sub-Chronic) Survival of *Mangifera indica* cold aqueous leaf extract treated

Flies

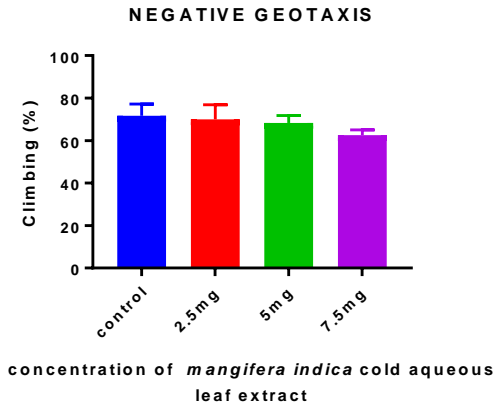


Fig 3: the climbing performance of *Mangifera indica* cold aqueous

Leaf extract-treated Flies.

3.5 Fecundity Assay Adult Flies or adult flies, Flies or flies

Paragraph The number of Adult Flies emerged from the egg laid by *Mangifera indica* cold aqueous leaf extract-treated Flies was used as a measure of fecundity. The number of emerged flies ranged from 7- 18. The highest number of emergence was recorded in the control while the least number was recorded in 10 mg/10 g diet. 17 and 11 emerged adult flies were recorded in 2.5 mg/10 g diet and 5 mg/10 g diet respectively. There was no significant difference ($p>0.05$) comparing the fecundity performance of the treated flies to the controlled flies. The result is presented in figure 4.

3.6 Acetylcholinesterase Activity

Paragraph Acetylcholinesterase activity of flies treated with *Mangifera indica* cold aqueous leaf extract ranged from 1.3-2.21 $\mu\text{mol}/\text{min}/\text{mgprotein}$. The highest activity was recorded in 2.5 mg/10 g diet and 5 mg/10 g diet while the least activity was recorded in the controlled flies. Acetylcholinesterase activity of 10 mg/10 g diet treated flies was recorded to be 1.94 $\mu\text{mol}/\text{min}/\text{mgprotein}$. There was no significant difference ($p>0.05$) comparing the

[fecundity performance](#) of the treated flies to the controlled flies. The result is presented in figure 5. [Fecundity](#)

[performance or Acetylcholinesterase activity?](#)

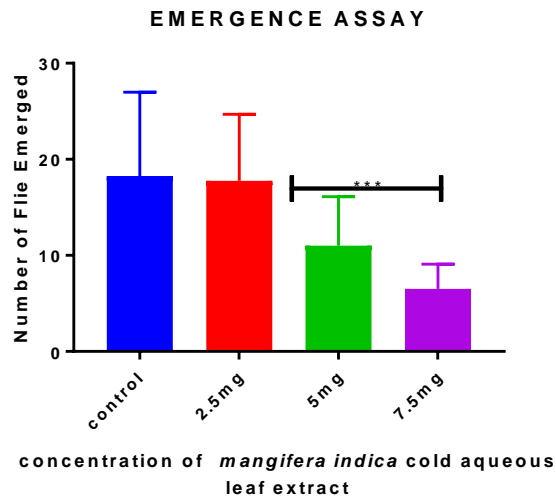


Fig 4: The number of emerged adult flies from the eggs of *Mangifera indica* cold aqueous

Leaf extract-treated flies.

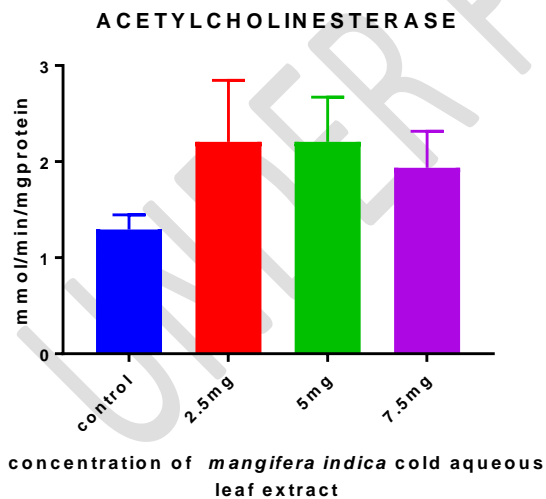


Fig 5: Acetylcholinesterase Activity of *Mangifera indica* cold aqueous extract treated flies

4.0 Discussion and Conclusion

4.1 Discussion

Why capital letter ?

Several **Processes** have been used for the isolation of phytochemicals from the natural products. Some of these processes are milling, grinding, homogenization and extraction. It has been reported that extraction is one of the important processes for separating phytochemicals from natural products [31]. The extraction effectiveness is affected by the chemical nature of phytochemicals, the extraction method used, sample particle size, the solvent used, as well as the presence of interfering substances [32]. It is important to depict the chemical nature of plant materials when their pharmacological activities are evaluated. Extracts revealed the presence of alkaloids, tannins, saponins, terpenoids, flavonoids and phenols as secondary metabolites. These secondary metabolites are reported to have many pharmacological properties, so this species is expected to have many medicinal uses [33]. Different alkaloids such as azadirachtin, azdirol, cymarol, digitoxin, kulactone, limocinin, salanin, toosendanin, xanthotoxin are contained in many plants etc which are soluble in water and toxic to herbivore insects. Many authors reported that the aqueous extract of plants has significant effect on insect mortality, their growth and reproduction [34-39]. From our result, the 100% acute concentration was found out to be 100 mg/10g diet while the 168 hrs LC₅₀ was calculated using probit analytical tool on Graphpad prism and it was found to be 72.4 mg/10 g diet. This might be due to some of these phytochemical components that are known to be toxic even at lowest concentration. The survival study (sub-chronic) of **flies** In our present study, shows high survival of controlled flies while there was gradual decrease in the survival of flies along concentration gradient (from 2.5 mg/10 g diet to 10 mg/10 g diet) and this might be traceable to some of these phytochemicals in *Mangifera indica* cold aqueous leaf extract. Fouad *et al.* [40], reported that the plant extracts possess broad spectrum toxic substances (toxic alkaloids, tannins etc.) that interrupt insect's normal physiology and influence on their feeding and mortality.

Negative geotaxis is the measure of how quickly a fly is able to climb vertically after being tapped to the bottom of a vessel as part of its innate escape response. Negative geotaxis is measured by either the distance an

animal is able to climb in a set time or the length of time it takes an animal to climb a set distance. Negative geotactic ability has been shown to be sensitive to oxidative stress, age, and cold exposure [41–44]. The geotactic ability of flies is impaired under abnormal conditions. From the result there was a slight decrease in the climbing activity of *Mangifera indica* cold aqueous leaf extract fed flies along concentration but statistically, there was no significant difference ($p>0.05$). This may be due to the alteration of the diet compositions with [mangifera indica](#) cold aqueous leaf extract. Linderman, *et al* [45]'s work showed a decline in negative geotactic ability in *L. monocytogenes* infected Oregon-R flies. Impaired climbing and flight behaviour in [Drosophila melanogaster](#) following carbon dioxide anaesthesia was a work carried out by Bartholomew, *et al.*, [46], who reported poor climbing performance of Flies at 5 min, 10min and 30 min CO_2 exposure. The climbing performance was worse with increase in exposure time or period. This negative geotactic ability has been useful in toxicological evaluation of toxic or poisonous plants, herbicide and neurotoxin.

Further, we sought to know if *Mangifera indica* cold aqueous leaf extract could affect the acetylcholinesterase (AChE) activity in the flies. Acetylcholinesterase is an enzyme that hydrolyzes acetylcholine at the post synaptic cleft. Acetylcholinesterase (AChE) is a key enzyme in the nervous system [47]. It terminates nerve impulses by catalyzing the hydrolysis of neurotransmitter acetylcholine. As a specific molecular target of organophosphate and carbamate pesticides, acetylcholinesterase activity and its inhibition have been early recognized to be a human biological marker of pesticide poisoning [47-49]. The success of this biomarker arises from the fact that it meets a number of characteristics necessary for the successful application of a biological response as biomarker in human bio-monitoring: the response is easy to measure, it shows a dose-dependent behavior to pollutant exposure, it is sensitive, and it exhibits a link to health adverse effects [47-51]. AChE activity determination is a sensitive marker of neurotoxicity, since inhibition of its activity can be an indicator of poor locomotor activity [52] and general toxicity [53]. From the result, we noticed significant increase in AChE activity which might have enhanced the climbing performance of the treated flies above 60% but there was no significant difference ($p>0.05$) compared to the controlled flies. Abolaji, *et al* [29] reported a contrary result of inhibited AChE activity of flies exposed to 4-vinylcyclohexene 1, 2-monoepoxide (VCM), and 4-vinylcyclohexene diepoxide (VCD). It has also been shown that the main active ingredient in cannabis, *tetrahydrocannabinol*, is a competitive inhibitor of acetylcholinesterase[54]. Similar to our result was reported by Agarwal *et al.*, [55], de Peyster *et al.*,

[56], a high activity of AChE and may inhibit the action of acetylcholine in the uterus to protect the pregnancy, thereby supporting fertility in female.

We sought to understand the impact of *Mangifera indica* cold aqueous leaf extract on flies fecundity and development. Reproductive fitness in insects has been well established. It has been reported that high activity of acetylcholinesterase plays important role in female fertility by catalyzing the hydrolysis of high concentration of acetylcholine in the Uterus protecting pregnancy [56]. In environments where nutrients are limited, there is a negative correlation between female fecundity and resistance to bacterial infection in *D. melanogaster* [57]. [immune](#) challenged females not only have fewer offspring, but those offspring also have shorter lifespan compared to the offspring of unchallenged female *D. melanogaster* [58] Reproductive fitness is a broad, ecological measure of health. Our result showed fewer offspring in the 5mg and 10mg *Mangifera indica* cold aqueous leaf extract-treated [Flies](#) compared to the controlled flies. In these 5 mg/10 g diet and 10 mg/10 g diet concentration some pupa did not [elcose](#) even after 12days of life cycle. Many authors reported that the aqueous extract of plants has significant effect on insect mortality, their growth and reproduction [34-39]. This may be due to oxidative stress induced by toxic phytochemical present in this extract at these concentrations.

4.2 Conclusion

Mangifera indica is commonly used in herbal medicine for one infection or the other and some authors have documented that it is one of the safe Medicinal plants with no record of LD/LC₅₀. From our study, we were able to determine the 168hrs LC₅₀ = 72.4 mg/10 g diet in *D. melanoogaster*. This may be due to the presence of alkaloids and tannins in the aqueous extract. The sub chronic result showed decrease in Flies survival along concentration with a least survival recorded in 10 mg/10 g diet. This might be due to stress induced from the consumption of the extract in their diet for a long period of time. The slight decrease and elevation of Climbing and AChE activities might be as a result of the effect of this extract consumption in flies. The delay in the development and eclosion of fewer flies in 5 mg and 10 mg/10 g diet might be as a result of the flies systems trying to adjust to the consumption of mangifera indica cold aqueous leaf extract in their diet. From this finding, *Mangifera indica* cold aqueous leaf extract might not be safe for long term consumption.

4.3 Recommendation

It is important to carry out the genetic study of *Drosophila melanogaster* fed diet with *Mangifera indica* aqueous leaf extract so as to draw a final conclusion on its toxicity in (*D. melanogaster*). I would change for “in this species.”

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