

LONG-TERM CONSUMPTION OF KOLA-NUT (*COLA NITIDA*) DIET DECREASED ANXIETY RELATED BEHAVIOUR IN MICE.

ABSTRACT

Following long-term consumption of kola nut (*cola nitida*) diet, anxiety related behaviour were studied in 16 Swiss white mice (18-28g body weight). The open field (OF) test, elevated plus maze (EPM) and the light/dark transition box (LD) tests were used. Swiss white mice were fed either control diet (rodent chow; n=8) or kolanut diet (50% w/w kola-nut diet; n=8) for 28 days. All animals were allowed free access to clean drinking water. Results showed that the frequency of rearing in the kolanut diet group was lower ($p < 0.05$) compared to control. The non-exploratory behaviours like grooming and genital licking were also lower in the test group compared to control ($p < 0.001$, 0.05 respectively). In the EPM test, the duration in the open arm in the kola diet group was higher compared to control ($p < 0.01$). The duration of grooming in the test group was however higher in the closed arm compared with control ($p < 0.01$). The frequency of downward dips only correlated positively with the duration in the open arm in the control [$r(16) = 0.855$; $p < 0.01$]. The kola fed animals spent more time in the light region of the LD test ($p < 0.01$) rearing and walling ($p < 0.05$), and spent less time in the dark region when compared with their control. In conclusion, long-term consumption of kola nut diet decreased anxiety-related behaviour in the mice.

Key words: Kola nut, anxiety

INTRODUCTION

Kola is a tropical tree crop (family - Sterculiaceae) with socio-economically important in Nigeria. Kola nut is used traditionally for ceremonies related to marriages, child naming, funerals and in other festivals and also chewed as a masticant (Ogutuga, 1975). It is commercially grown in the West where it is known as "Obi" in Yoruba, consumed by the Northerners where it is known as "Goro" in Hausa, and revered in the East where it is called "Oji" in Igbo. In Cross River and Akwa Ibom it is called "Ibong" in Efik. It is of great importance in the traditional institution hence the adage by the Igbos ... "He who brings Kola brings life". Kola is generally ascribed to elders' privileges.

The nuts are either chewed whole or used in powdered state. It is known to cause mild stimulation of the central nervous system and produce a temporal feeling of increased physical strength often associated with a reduction of hunger and fatigue, which may be due to its high content of carbohydrates (Martin et al, 1983). It was reported by Egbe and Oladokun (1987) that there are over forty kola species out of which *cola nitida* and *cola acuminata* are of major economic importance in Nigeria. Fresh and cured kola nut chewed in small doses increase mental activity, reduce the need to sleep and also dispel hunger and thirst (Ogutuga, 1975). It is for

this reason that kola nut chewing has become very popular among students, drivers and many other consumers who need to remain active for unusually long periods.

In some developed countries, however, kola nut extracts are used industrially for the manufacturing of many cola-type soft drinks flavours (Beattle, 1970), as a source of caffeine used for many manufacture of pharmaceutical products and essential oils (Olounloyo, 1979) and as a main ingredient in production of heat-tolerant chocolate bars (Williams 1979). In addition, caffeine is known to be a fat burner and therefore beneficial in assisting weight loss (Blades, 2000). As a result of the commercial importance of kola nuts, a lot of research work has been done on Cola Nitida, the kola of commerce, in Nigeria (Oladokun, 1982). Presently, the bulk of kola nuts being produced in Nigeria are either consumed fresh locally or exported as sun-dried to some drier areas of Africa, where they are used as masticant or as sources of colourant for cloth dyeing but with little or no industrial use locally (Jayeal, 2001).

Since kola nut contains caffeine which has the stimulating effect on the nervous system, it is likely that chronic consumption of kola nut will have an effect on anxiety-related behaviour. Therefore the effect of long-term consumption of kola nut on locomotor activities and anxiety-related behaviour in mice was studied using the open field test, elevated plus maze test and light dark transition test.

MATERIALS AND METHODS

Swiss white mice purchased from the animal house of the Department of Physiology, University of Calabar. Mice were grouped into two: Control (n=8) given normal rodent chow (Vital feed Nigeria), and test (n=8) given 50% w/w kola-nut diet. Kola-nut diet was prepared by slicing, drying and grinding fresh kola-nut (*colc nitida*) bought from Bogobiri (Hausa Market) in Calabar, Nigeria. Equal portions of the grinded kola-nut powder (10g) and rodent chow (10g) were used to constitute 50% w/w kola-nut diet (Osim and Udia, 1993). All animals had free access to clean drinking water. This feeding was done for 28 days. All animals were weighed before and after the feeding period.

The **open field maze** test used by Bisong *et al* (2018) was employed in this study. The test apparatus measured 72 x 72 x 32cm (l x b x h) with a floor divided into sixteen 18 x 18cm squares, and centre square of 36 x 36cm. Mice were allowed 5 minutes to explore the apparatus while behaviour are scored. (Brown *et al.* 1999).

The **elevated plus maze** as described by Lister (1987) and used by Bisong *et al* (2017, 2018) was employed in this study. The apparatus has 2 open arms (each measuring 30x5x15cm) and two closed arms (each measuring 30x5x15cm) extending from a central, open square (5x5cm). The maze was elevated on a pedestal to a height of 45cm above the floor. Each mouse was placed in the centre square of the elevated plus maze facing an open arm and its behaviour scored.

The **light/dark transition box** test as described by Hascoët and Bourini, (1998) and used by Bisong *et al* (2017) was used. It is box measuring 46x30x27cm high (l x b x h cm), divided into two compartments; a small 18x30cm area and a large 27x30cm area with a 7x7cm door on floor of the partition linking the two chambers. The small compartment was painted black to mimic darknes, whereas the large compartment was painted white. Mice were given 5 minutes to explore the apparatus during which behaviours were scored.

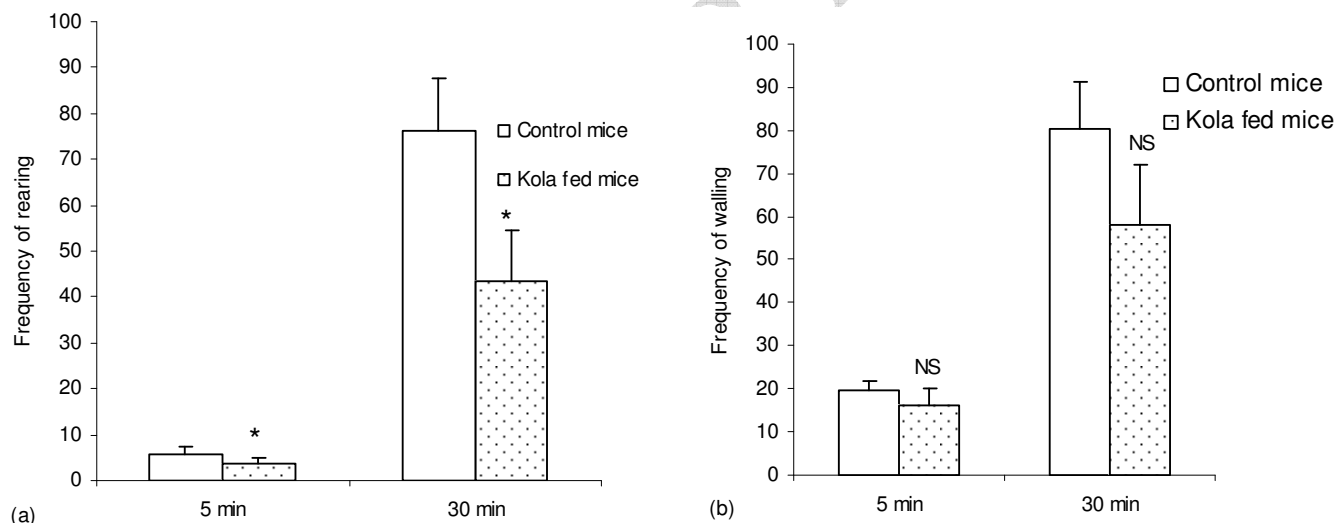
Statistical Analysis

Data obtained from the study were analysed using the Student T- test. Associations between data were tested using the Pearson's correlation. Data were presented as means \pm standard error of mean. Probability level of $P < 0.05$ was accepted as significant.

RESULTS

Exploratory behaviour in the open field maze

The Exploratory behaviour, Rearing and walling, are forms of vertical locomotor activity. The frequency of Rearing in the test group within the first five minutes was 3.6 ± 1.1 , significantly lower when compared with control (5.9 ± 1.5 ; $P < 0.05$). At the end of 30 minutes, the frequency of rearing in the kola group (43.4 ± 11.1) was also lower than that in control (76.4 ± 11.2 ; $P < 0.05$). Although the frequency of walling in the kola group was lower both at 5 minutes (16.4 ± 3.6) and 30 minutes (58.3 ± 14.0) when compared with control (19.6 ± 2.3 at 5 minutes and 80.4 ± 10.7 at 30 minutes; Figure 1).



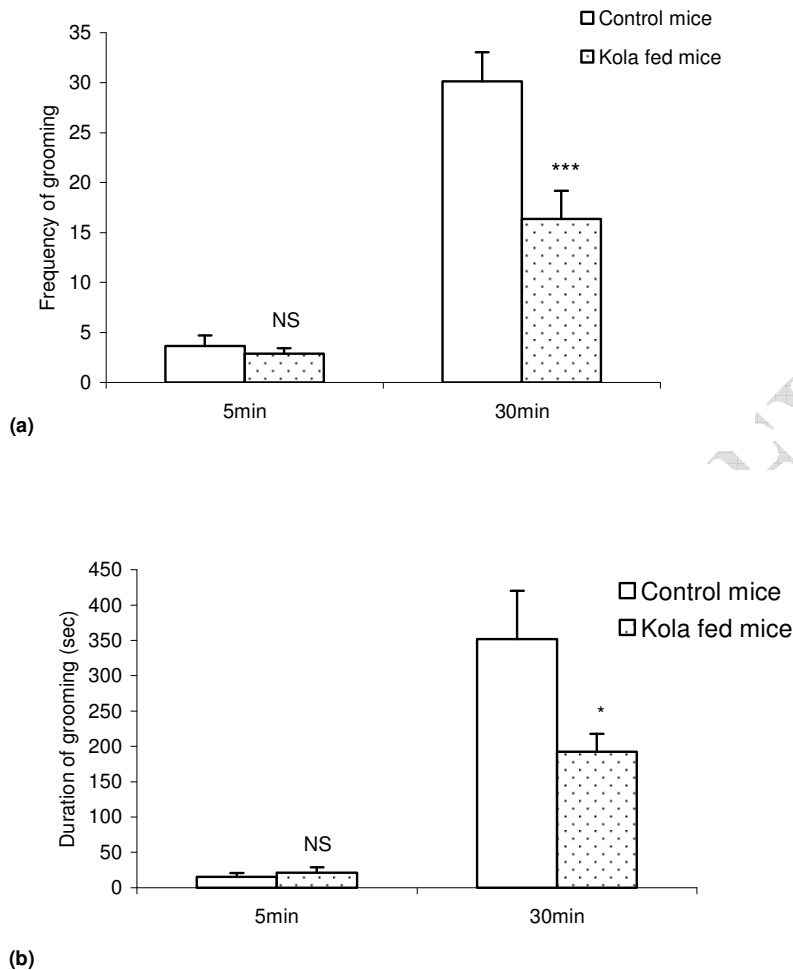
NS – Not significant compared to control
* – Significant at $P < 0.05$ compared to control

Fig. 1: Exploratory behaviour; rearing (a) and walling (b) of mice fed kola diet in the open field maze.

Grooming in the open fields maze

The frequency and duration of grooming in the open field maze were significantly lower in the test group than in the control at the end of the 30 minutes session. This was $16.4 \pm 2.8/30\text{min}$ in the kola group, which was less than that in the control group of animals, which was 30.1 ± 2.9 ; ($P < 0.01$). The duration of the

150 grooming at the end of 30minutes (192.5 ± 25.4 sec) was also significantly lower in
 151 the kola fed group compared to control (352 ± 54 sec; $P < 0.05$). This is shown in
 152 Figure 2.
 153



154

155

156

157

158

159

160

161

162

163

164

165

166

167

168

169

170

171

172

NS – Not significant compared to control
 * - Significant at $P < 0.05$ compared to control.

Fig. 2: Frequency (a) and Duration(b) of grooming in mice fed kola diet in five minutes and thirty minutes in the in the open field maze.

Comparison of activities in the Elevated plus maze between kola fed and control mice.

The frequency of entry into the open arm was not significantly different between control and kola fed mice. The duration in the open arm was however, significantly higher in the test group compared to control ($p < 0.01$). The frequency and duration of downward dipping of mice did not differ significantly between test and control groups of mice. The frequency of entry into the closed arm of the elevated plus maze in the kola fed group was significantly lower compared to control ($P < 0.05$). The duration in the closed arm was also lower in the test group compared

173 to control ($P < 0.05$). Although the frequency of grooming in the closed arm was not
 174 significantly different between test and control groups, the duration differed ($P < 0.01$).
 175 The frequency of rearing and the number of faecal boli at the end of the 5minutes
 176 session did not differ from control values. Table 1 below shows a summary of these
 177 comparisons.

178
 179
 180
 181
 182
 183

Table 1: Summary of comparison of activities in the Elevated plus maze between kola fed and control mice.

Parameters	Control group	Kola fed group	Level of significance
Frequency of open arm entry (/5minutes)	2.8 ± 0.3	2.6 ± 0.6	NS
Duration in open arm (seconds)	45.3 ± 10.4	136 ± 18.7	0.01
Frequency of downward head dipping (/5minutes)	11.4 ± 2.9	12 ± 2.2	NS
Duration in closed arm (seconds)	218.1 ± 15.2	158.9 ± 25.5	0.05
Frequency of grooming in the closed arm (/5minutes)	3.3 ± 0.8	4 ± 0.5	NS
Duration of grooming in the closed arm (Seconds)	18.3 ± 5.7	44.9 ± 9.3	0.01
Frequency of genital licking in the closed arm (/5 minutes)	1.3 ± 0.4	2.3 ± 0.5	0.05
Frequency of rearing in the closed arm	19 ± 2.0	20 ± 2.1	NS
Number of faecal boli	0.4 ± 0.3	0.3 ± 0.2	NS

184 NS – Not significantly different compared to control.

185
 186
 187
 188
 189
 190
 191
 192
 193
 194
 195
 196

Correlation between duration in the open arm and frequency of downward head dipping

There was a positive correlation between the duration in the open arm of the elevated plus maze and the frequency of downward head dipping in the control group [$r(16) = 0.855$; $p < 0.01$], Figure 3.

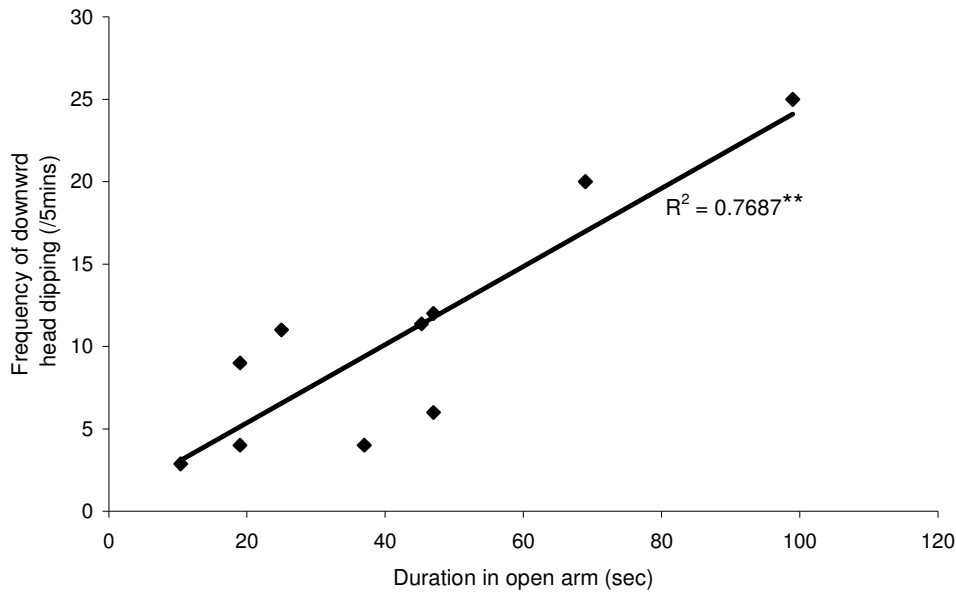


Fig. 3: Correlation between frequency of downward head dipping and duration in the open arm in the control group

The Effect of long-term feeding with kola diet on Activities in the light region of the light/ dark transition box.

Activities in the light region were generally higher for the kola fed group compared to control. Although the frequency of entry into the light region did not differ, the time spent (duration) in the light region was significantly higher in the test group (111.5 ± 9.43 seconds) compared to control (77.63 ± 12.7 seconds; $p < 0.05$). The frequency of line crossing did not differ between the test and control groups. The frequency of rearing and walling were higher in the test groups compared to control ($p < 0.01$; $p < 0.05$ respectively), Table 2.

Table 2: Activities in the light region of the light/ dark transition box.

Activity in the light region.	Control group	Kola group	fed	Level of significance
Frequency of entry (/5mins)	6.75 ± 0.90	6.0 ± 0.95		NS
Duration in light region (Sec)	77.63 ± 12.7	111.5 ± 9.43		0.05
Line crossing (/5mins)	59.0 ± 10.29	51.3 ± 6.43		NS
Frequency of rearing (/5mins)	3.13 ± 0.82	9.25 ± 1.65		0.01
Frequency of walling (/5mins)	7.75 ± 1.75	14.88 ± 2.76		0.05

NS – not significant compared to control

The Effect of chronic feeding with kola diet on Activities in the dark region of the light/ dark transition box.

Although the test group showed less activity in the dark region of the light/dark transition box, the exploratory activities were significantly higher compared to control. The frequency of entry and the time spent (duration) in the dark region did not differ in the test group when compared to control. Frequency of rearing in the test group was higher than the value for the control group ($p < 0.05$). Walling also followed a similar trend with the test group of animals walling more than control ($p < 0.05$). The frequency of grooming did not differ significantly but the duration of grooming was lower in the test group compared to control ($p < 0.05$). The frequency of genital licking in the test group of mice was also significantly lower compared to that in the control group ($p < 0.05$); Table 3.

Table 3: Activities in the dark region of the light/ dark transition box.

Activity in the dark region.	Control group	Kola fed group	Level of significance
Frequency of entry (/5mins)	7.75 \pm 0.90	5.13 \pm 0.83	NS
Duration in dark region (Sec)	222.5 \pm 12.37	197 \pm 13.97	NS
Frequency of rearing (/5mins)	8.5 \pm 1.56	14.0 \pm 3.1	0.05
Frequency of walling (/5mins)	12.88 \pm 2.53	20.4 \pm 2.74	0.05
Frequency of grooming (/5mins)	6.38 \pm 6.0	6.0 \pm 1.12	NS
Duration of grooming (sec)	63.88 \pm 13.73	32.6 \pm 6.43	0.05
Frequency of genital licking (/5mins)	4.5 \pm 1.09	2.0 \pm 0.61	0.05

NS – not significant compared to control

DISCUSSION AND CONCLUSION

The open field maze has been used to assess the emotionality of animals in a novel environment, as well as locomotion and exploration (Weiss and Greenberg, 1996). Although the frequency of line crossing did not differ significantly between the test and control groups, other forms of locomotor activity differed. This implies that there was no significant change in horizontal locomotor patterns.

The frequency of rearing was significantly lower in the kola fed group (test) when compared to control in the open field. This trend was also similar in the frequency of walling. The vertical locomotor activities and therefore exploratory activity were decreased following long-term ingestion of kola nut. The frequency and duration of grooming in the kola fed group was also significantly decreased at the end of 30 minutes. This implies decreased vertical locomotor (exploratory) activities following chronic ingestion of kola nut. Therefore long-term ingestion of 50%w/w kola nut diet decreased exploratory activity. It is most likely that consumption of large

quantities of cola nut will not serve the stimulant effect any longer but rather cause depression of the nervous system. These results however, do not support the report of previous researchers that kola nuts serve as a stimulant (Ogutuga, 1975; Martins 1983). The decrease in exploratory/vertical locomotor activity following long-term consumption of kola is in consonance with reports of Neil (1978) which showed that excessive consumption of caffeine caused mixed depressive states in psychiatric patients; and also the work of Greden (1978) depressive syndrome as being associated with caffeine, which is one of the major constituents of kola nut.

The elevated plus maze has been proven as a model for assessing anxiety and fear (Brown et al, 1999, Lister, 1987). This test is based on the natural aversion of rodents for open space and heights. Mice fed 50%w/w kola diet when compared with control, spent more time in the open arm and less time in the closed arm. These therefore means that the kola fed mice were less fearful compared to their control. To buttress these, there was a positive correction between the duration in open arm and the frequency of downward head dipping. Therefore, the less fearful mice spent more time in the open arm of the maze and did more of the downward head dips.

The light/dark transition box is also used as a model for assessing anxiety and fear. This light/dark test is based on the innate aversion of rodents for brightly illuminated areas and on the spontaneous exploratory behaviour of rodents in response to mild stressors that is novel environment and light (Crawley and Godwin, 1980). In this test, the kola fed group of mice spent more time in the illuminated (light) region of the and also showed increased activity (rearing and walling) in this region. This implies that long-term feeding of mice with kola diet produces an anxiolytic effect.

The kola fed group showed less grooming (non-exploratory behaviours associated with fear) in the dark region compared to their control. The frequency of the exploratory behaviours, rearing and walling, were higher in the test group. This is in agreement with findings of Costal *et al* (1989) that increased exploratory behaviours was associated with an increase in the time spent in the light region of the light/ dark box.

The mice in the test group were less fearful compared to control, which is in consonance with the test in the elevated plus maze. Therefore, long-term ingestion kola diet caused an anxiolytic effect. These results were contrary to earlier works which implicate caffeine (a major component of kola) as an anxiogenic agent (File and Hyde, 1979; Charney *et al*, 1984).

In conclusion, long-term consumption of 50%w/w kola diet in mice caused decreased locomotor activity, instead of producing a stimulant effect, in mice. The kola diet also produced an anxiolytic effect, thereby reducing fear and anxiety in mice.

REFERENCES

- Acher, J (1973): Tests for Emotionality in Rats and Mice: A Review. *Anim. Behav.* 21: 205 – 235
- Beattle, G.B (1970): Soft drink flavours. Their History and Characteristics. 1 Cola or "Kola" flavours the flavour Industry pp. 390 – 394

301 Bisong S. A., U. E. Okon, E. A. Egbung, F. E. Abuo¹ and O. A. Sanya (2017). Effect of
 302 Crude Ethanol Leaf-extract of *Murraya koenigii* on Anxiety in Mice. *Asian*
 303 *Journal of Medicine and Health* 7(1): 1-9; Article no.AJMAH.36294

304 Bisong S. A., C. O. Nku K. U. Nwoke and E. E. Osim (2018). Crude aqueous leave
 305 extract of *Carica papaya* Linn (pawpaw) reduced anxiety and fear related
 306 behaviour in CD1 mice. *European Journal of Pharmaceutical and Medical*
 307 *Research*, 2018, 5(3), 488-493.

308 Blades, M (2000): Functional foods or Nutraceutical Nutrition and food science vol.
 309 30 Number 2 pp 73 – 75

310 Brown, R.E, Corey, S.C; Moore, A.K (1999): Differences in Measures of Exploration
 311 and Fear in MHC – Congenic C57BL/6J and B6 – H – 2k Mice. *Behaviour*
 312 *and Genetics*; 26, 263 – 271.

313 Charney, D.S; Galloway, M.P; Heninger, G.R (1984): The Effects of Caffeine of
 314 Plasma MHPG; Subjective Anxiety, Automatic Symptoms and Blood
 315 Pressure in Healthy Human. *Life Sci.* 35: 135 – 144

316 Crawley, J. N., Godwin, F.K. (1980). Preliminary report of a simple animal
 317 behaviour for the anxiolytic effects of benzodiazepines.
 318 *Pharmacol. Biochem. Behav.* B, 167 – 170

319 Egbe, N.E and Oladokun, M.A.O (1987): Factors limiting high yield in kola nut (*cola*
 320 *nitida*) production Nigeria. *Café cocoa*, the vol. XXXL, 1004, Oct. - Dec.

321 File, S.E; Hyde, J.R.G (1979): A Test of Anxiety that Distinguishes Between the
 322 Actions of Benzodiazepines and those of other minor Tranquilizers and of
 323 Stimulants. *Pharmacol. Biochem. Behav.* 11: 65 – 69

324 Greden, J.F *et al* (1978): Anxiety and Depression Associated with Caffeine Among
 325 Psychiatric patients. *Am. J* 131, 8, 963

326

327 Hascoët, M; Bourin, M (1998): A New Approval to the Light/Dark Procedure in Mice.
 328 *Pharmacological Biochem. Behav* 60, 645 – 653

329 Jayeal, C O (2001): Preliminary studies on the uses of kola nuts (*cola nitida*) for soft
 330 drink production. *The journal of food Technology in Africa*, Vol. 6, No. 1,
 331 pp. 25 – 26

332 Lister, R.G. (1987): The Use of a Plus-Maze to Measure Anxiety in Mouse.
 333 *Psychopharmacology* 92: 180 – 185

334 Martin, K.L; Morelli H. F, Schild, H.O; State land, B.E (1983): In *Clinical Pharmacology*
 335 *Basic Principles in Therapeutics*, 2nd (Ed.) Macmillan Publishing Co. Inc.
 336 New York p. 663

337 Neil, J.F (1978): Caffeinism Complicating Hypersomnic Depressives Episodes.
 338 *Comprehensive Psychiatry* 19, 377

339 Ogotuga, D.B.A. (1975): Chemical composition and potential commercial uses of Kola
 340 nuts, *Cola nitida* vent cachott and Endlisher. *Ghana J-Agric Sci.* 8, 121-
 341 125

342 Oladokun, M.A.O (1982): Morph-physiological aspect of germination, rooting and
 343 seedling growth in kola (*cola spp*) Ph. D thesis, the University of Ibadan
 344 pp. 230

345 Olounloyo, O.A (1979): Fungi Associated with deterioration of high kola nuts. *Nigerian*
 346 *Journal of Agric Science* (1) 52-59

347 Osim, E.E and Udia, P.M (1993): The effect of consuming kola nut *cola nitida* diet or
 348 mean arterial pressure in rats. *Int. J. pharmacog.* 31, No3, 193 - 197

349 Rodgers, R.J (1997): Animal Models of “Anxiety”: Where Next? *Behav. Pharmacol.* 8:
 350 477 – 496

- 351 Streng, J (1974): Exploration and Learning Behaviours in Mice Selectively Bred from
352 High and Low Levels of Activity. *Behav. Genet.* 4: 191 – 204
- 353 Weiss, G and Greenberg, G. (1996). *Open field procedures*. In Greenberg, G. and
354 Haraway, M.H. (Eds) *Comparative psychology. A hand book*, Gaillard
355 New York. pp. 603-62.
- 356 Williams, S.O (1979): Project of kola chocolate processing and consumption CRIN,
357 (Seminar paper 4).

UNDER PEER REVIEW