TOOTHPASTE (CLOSE UP) IN MALE ALBINO

RATS (*Rattus norvegicus*)

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8 Abstract

9 This study was carried out to determine the biomarker response of male albino wistar rats (Rattus novergicus) to a daily dosage of toothpaste. Twenty four wistar rats were divided 10 randomly into two groups and housed in wooden cages. The first group which is the test 11 group was administered with varying doses (250ul, 270ul, 300ul) according to their body 12 weight (0.00167mg/g body weight) per week for three weeks while on the fourth week no 13 treatment was given. This was done to observe the rate of recuperation from effects of 14 treatment. The second group which was the control group were given distilled water of equal 15 16 measurement with the treatment given to the test rats. Several biochemical and hematologic 17 parameters were used to evaluate the effect of toothpaste. Parameters used were; for enzyme 18 and liver functions, alkaline aminotransferase (ALT), aspartate aminotransferase (AST), and protein, for kidney sodium (Na+), potassium (K+), chloride (Cl) and bicarbonate (HCO₃) 19 while for hematology white blood cells (WBC), red blood cells (RBC), platelets, lymphocytes, 20 hemoglobin and packed cell volume (PCV) and sperm count was also used. The results 21 showed significant difference (P < 0.05) in the parameters when compared with the control 22 group. These findings demonstrate that toothpaste caused detrimental effect on sperm 23 parameters which could lead to infertility in males. There were also observed changes in 24 liver, blood parameters and kidney which could lead to renal dysfunction when exposed to 25 this substance for extended periods. The intentional or accidental ingestion of toothpaste 26 27 should be avoided especially in children.

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30 1.0 INTRODUCTION

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Toothpaste is a personal care product used by millions across the world. Despite the different brands, they all have some major active ingredients that are general to all of them and essential in making toothpaste. These ingredients are fluoride (sodium fluoride), abrasives (hydrated silica) and detergents (sodium lauryl sulphate, SLS). Other inactive ingredients present are flavor, sorbitol etc. (ADA, 2017). Sodium fluoride being a major component of toothpaste is an inorganic salt. It is a chemical compound and an odourless, colourless crystalline solid (Spellman, 2008) that came into use to prevent tooth decay in the 1940s 39 (Murray et al., 2003). It has a molecular formula of NaF. It is white to greenish in colour 40 depending on its level of purity (Haynes, 2011; British Medical Association., 2015). It is non-41 combustible and corrosive to aluminium metal, it is known to be insoluble in alcohol but 42 highly soluble in water (O'Neil, 2001). Sodium Fluoride is used not just as fluorinate in 43 toothpaste but also in the preservation of wood, as a corrosion inhibitor, insecticide, cleaning 44 agent, chemical reagent and in glass and metallurgy industries (Aiguesperse et al., 2005). 45 Fluoride has been studied extensively for use in the medical industry (Haguenauer et al., 2000). Sodium fluoride is generally safe for dental health at low concentrations but 46 47 continuous ingestion of large amounts of sodium fluoride poses possible dangers to health, 48 with short term exposures causing irritations to eyes, skin and nasal membranes (Green, 49 2005). Studies have shown that fluorides, especially when in solution forms (aqueous forms) 50 are more extensively absorbed into the body and are classed as toxic by both inhalation and 51 ingestion through oral routes (Kapp, 2005) The rate at which fluoride (as Sodium Fluoride) is 52 absorbed is inversely related to the pH of the stomach contents (WHO, 2006). Acute 53 exposure and toxicity can result in nausea, abdominal pain, and diarrhea. Other possible 54 effects are muscle paralysis, extremity spasms (Whitford, 2011). Study has shown that 55 continuous ingestion of fluoride causes deleterious effects on skeletal (Cheng *et al.*, 2008), 56 dental (Flaitz et al., 2000), soft tissues (brain), thyroid (Bathnagar et al., 2005) and testis 57 (Wan et al., 2006). In a study it was observed and documented by Shashi, (2003) that 58 fluoride exposure can induce the loss of neuronal cell bodies and damage synaptic structures 59 in different regions of the brain (Gopalakrishna et al., 2002) as well as cause inhibition of 60 enzyme activity and a decrease in expression of membrane proteins (Barbar et al., 2006). In 61 the blood and liver of animals it was observed that various changes like abnormal behavioural 62 patterns and metabolism occur after chronic administration of fluoride lesions (Ramakrishna 63 and Saralakumari, 1991; Denbesten et al., 1995).

64 Beyond Sodium Fluoride, Sodium lauryl Sulfate (SLS) is also another major constituent of toothpaste; Sodium lauryl sulfate (SLS), also known as sodium dodecyl sulfate, is an anionic 65 66 surfactant commonly used as an emulsifying cleaning agent in household cleaning products 67 (laundry detergents, spray cleaners, and dishwasher detergents) (Cara et al., 2015), it's low 68 cost and desirable action as a foaming agent has led to its use in the formulations of 69 toothpaste (Lippert, 2013). Like all detergents, SLS has been shown to cause skin and eye 70 irritation and cause more skin related damage especially with prolonged exposure (Cara, et al., 2015). A research carried out by Cosmetic Ingredient Review (2015) on the health and 71 72 safety of the SLS chemical using rats as test subjects showed that SLS is harmful by the oral

73 route, while using rabbits and guinea pigs as test subjects it was found to be harmful in the 74 dermal route. SLS was also reported to irritate the respiratory tract and cause irritation in both 75 skin and eye of rabbits. No gross lesions or microscopic abnormalities were found in a chronic oral feeding study in rats given 0.25%, 0.5% and 1.0% of SLS in their diet for two 76 77 years (Fitzhugh and Nelson, 1968) and the same result was observed in using a different test 78 subject in a chronic oral one-year oral toxicity study using beagle pups with 0%, 0.67%, 79 1.0%, or 2.0% SLS. This study is aimed at evaluating the possible effects of toothpaste 80 ingestion (accidentally or intentionally) on hepato-renal functions, hematological and sperm 81 parameters in male albino rats and associating such effect on humans.

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2.0 MATERIALS AND METHODS

84 2.1 *Experimental setup*

24 albino wistar rats (Rattus norvegicus) were used. The animals were weighed and randomly 85 86 allocated into two experimental groups ose Up toothpaste, a popular brand of toothpaste 87 used here in Nigeria was administered to the rats in mimicking concentrations commonly 88 used daily. 1ml of the toothpaste was dissolved in 100ml distilled water to make a solution. 89 The estimated average daily human dosage of toothpaste used was calculated and measured 90 and the same dosage was administered to the rats. They were calculated using the weights of 91 the rats and the dosage administered ranged depending on the change of the weekly body 92 weights of the rats. The oral rout was used for administration, using a 1ml syringe. The 93 experiment was carried out for four (4) weeks. The treatment was administered to the test 94 group for three weeks while on the fourth week no treatment was given to the test group. This 95 was done to observe how their body adapts and tries to recuperate and handle the effects from 96 the treatment substance. Three (3) rats from the test group were sacrificed weekly. While 97 three (3) from the control group were sacrificed weekly. This was done to enable us collect 98 blood and sperm samples for analysis and to allow for careful observation of the specific 99 organs of the rats. Before each sacrifice each rat was weighed and its final body weight was 100 recorded after overnight starvation. The animals were sacrificed by jugular puncture while 101 under chloroform anaesthesia Bood samples collected were taken with both EDTA and 102 Heparin bottles for laboratory analysis while the testes were collected for sperm analysis 103 which was done using an electron microscope.

105 2.2 Biochemical Analysis

106 Standard procedures were ensured during the collection of the blood, sperm and liver samples 107 prior to biochemical analysis. The plasma activity of Alkaline Phosphatase (ALP) was determined using Radox kit (colorimetric method) of Rec (1972). Biuret method was used to 108 109 determine the level of total protein in the samples according to the method of Flack and 110 Woollen (Flack and Woollen, 1984). The plasma activity of aspartate transaminase was 111 determined using Reitman and Frankel method (Reitman and Frankel, 1957). The serum 112 electrolytes were determined using ISO 4000 Automated electrolyte analyser. SFRI, France. 113 The plasma activity of alanine transaminase was determined using Reitman and Frankel 114 method (Reitman and Frankel, 1957). The sperm motility, viability and abnormalities were 115 determined using one step eosin method and the epididymal sperm count was done with 116 Neubauer haemocytometer (Deep 1/10 mm, LABART, Munich, Germany) and light 117 microscope at $40 \times$ magnifications.

118 2.3 Data Analysis

119 Data were analyzed using Tukey test at a level of 5% probability, using Assitat Software

120 Version 7.7 en (2017).

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122 **3.0 RESULTS**

123 The effects of oral administration of Close Up toothpaste on the Hepato-renal parameters in 124 male albino rats are presented in Table 3.1. The result showed significant difference in the 125 levels of electrolytes and hepatocyte parameters between the Test and Control across each 126 week and between the Test and average control (four week) in each week. Results from the 127 first week revealed a higher value of sodium (Na) on test compared to the control with a significant difference (P < 0.05) but no significant difference (P > 0.05) among the test of 128 129 potassium (K), chlorine (Cl), ALT, AST and their respective control. On the second week, 130 there was no significant difference (P > 0.05) among the test and the respective controls of 131 sodium (Na), potassium (K), bicarbonate, AST and ALT. While on the third week, the 132 analyzed result showed non-significant difference (P > 0.05) among sodium (Na), potassium

(K), bicarbonate, AST and ALT and their respective control, except chlorine (Cl), which 133 134 showed a significant difference (P < 0.05). Finally, on the fourth week, the result showed that there was significant difference (P< 0.05) among sodium (Na), potassium (K), chlorine (Cl), 135 136 bicarbonate, ALT, AST and their respective control. The result also showed the various significant differences between the Test and the average control. The result on Sodium 137 138 showed no significant difference between week one, week two, week three against the 139 average control but showed significant difference (P>0.05) in week four. The result on 140 Potassium (K) showed no significant difference between week one, week two against the 141 average control at (P>0.05) but shows significant difference (P< 0.05) between the tests of 142 week 3 and week 4. The result on chlorine (Cl) revealed there were no significant difference 143 (P>0.05) between week one, week two, week three against the average control, but there 144 were significant difference (P < 0.05) in the fourth week. The result on bicarbonate showed 145 there were no significant difference (P>0.05) between week one, week two, week three, week 146 four and the average control. The result on ALT, showed significant difference between week 147 one, week two, week three, week four and the average control at (P < 0.05). Finally, the result 148 on AST showed significant difference between week one, week two, week three, week four 149 and the average control at (P < 0.05). 150

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TABLE 3.1: RESULT SHOWING THE EFFECT OF TOOTHPASTE ON SODIUM, POTASSIUM, CHLORIDE, BICARBONATE,
 AST AND ALT

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		Na (mmol/L)	K (mmol/L)	Cl (mmol/L)	HCO3 (mmol/L)	AST (UI/L)	ALT (UI/L)
	Control	133.67 ± 2.50^{a}	4.05 ± 0.25^a	100.67 ± 4.5^{a}	23.67 ± 0.5^{a}	17.67 ± 3.50^{a}	10.67 ± 1.50^{a}
WEEK 1	Test	$143.00 \pm 4.00^{b,A}$	$3.60\pm0.20^{a,AB}$	$99 \pm 1.00^{a,A}$	$23\pm0.00^{a,AB}$	$27.33 \pm 8.50^{a,B}$	$11.67 \pm 0.50^{a,B}$
	Control	157.67 ± 22.50^{a}	7.25 ± 2.55^a	109.67 ± 18.50^{a}	23.67 ± 1.50^a	34.67 ± 3.50^a	10.0 ± 2.00^a
WEEK 2	Test	$138.67 \pm 12.50^{a,A}$	$4.38\pm0.05^{a,AB}$	$95\pm7.00^{a,A}$	$25\pm4.00^{a,A}$	$29.67\pm1.50^{a,AB}$	$6.67\pm0.50^{a,C}$
	Control	136.67 ± 10.50^{a}	5.0 ± 0.60^{a}	86.67 ± 4.50^a	24.67 ± 3.50^a	23.67 ± 5.50^a	11.0 ± 4.0^a
WEEK 3	Test	$129.0\pm1.00^{a,\mathrm{AB}}$	$3.9\pm0.30^{b,AB}$	$85\pm1.00^{a,ab}$	$19.67\pm0.50^{a,B}$	$30.33\pm3.51^{a,\mathrm{AB}}$	$12.67\pm0.5^{a,B}$
	Control	149.67 ± 0.50^{a}	5.10 ± 0.10^a	106 ± 1.00^a	23.0 ± 1.00^a	23.0 ± 1.00^{b}	13.06 ± 1.0^{b}
WEEK 4	Test	$111.67 \pm 3.50^{b,B}$	$2.9\pm0.20^{\text{b},B}$	$76.66\pm4.50^{b.B}$	$20.0\pm1.00^{\text{b},\text{AB}}$	$45.0\pm4.00^{a,A}$	$24.67 \pm 1.5^{a,A}$
AVERAGE CONTROL	Control	142.50 ± 11.83^{A}	$5.43 \pm 1.13^{\rm A}$	98.83 ± 9.16^{A}	23.83 ± 1.83^{AB}	25.16 ± 4.16^{B}	10.50 ± 2.5^{B}

163 ^{a-b}Different letters in the same column indicate significant difference (P<0.05) within each week

^{A-B} Different letters in the same column indicate significance difference (P<0.05) across the weeks

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167 For the haematological parameters, on the first week, there was a higher value on the test 168 samples compared to the control in the Packed Cell Volume (PCV), haemoglobin (Hb) and red blood cell (RBC) with a significant difference (P < 0.05) between the test and control 169 170 while for White blood cells (WBC), Platelets and Lymphocytes showed non-significant differences (P > 0.05) between the test and average. The second and third week both showed 171 172 no significant difference (P> 0.05) among Packed Cell Volume(PCV), Haemoglobin(Hb) and 173 Red blood cell (RBC), White blood cells (WBC), Platelets while Lymphocytes showed a 174 significant difference (P < 0.05) between the test and control. In the fourth week there were 175 significant difference in all hematological parameters except Red blood cells (RBC).

176 There were no significant difference (P> 0.05) in Packed cell volume (PCV) in week1, 177 week2, week3 when compared with the average control but there was a significant difference 178 (P < 0.05) in the week 4. No significant difference (P > 0.05) was seen in the fourth week for 179 Haemoglobin between the test and average control but significant difference (P < 0.05) was 180 noted all through the first three weeks. No significant difference was seen in both Red Blood 181 Cells (RBC) and White blood cells (WBC) through the four weeks when the test was 182 compared with the average control. Platelets showed significant difference (P < 0.05) across 183 all four weeks when the test and average control were compared. Lymphocytes showed no 184 significant difference all through the four weeks when the test is compared to the average control. In the result for semen analysis, results from week 1 to week 4 all had a lesser value 185 186 of sperm count on the test when compared to the control, with a significant difference 187 (P < 0.05) between the control and the treatment although the result showed no significant 188 difference (P > 0.05) between the test and the average control across the four weeks.

TABLE 3.2: RESULT OF THE EFFECT OF TOOTHPASTE ON PROTEIN, PACKED CELL VOLUME, HEMOGLOBIN, RED BLOOD CELLS, WHITE BLOOD CELLS, PLATELETS, LYMPHOCYTES

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		PROTEIN (g/L)	PCV (%)	Hb (g/dl)	RBC (X ¹²)	WBC (X ⁹)	PLATELETS (X ⁹)	LYMPHOCY TES (X ⁹)
	Control	67.70 ± 12.19^{a}	26.50 ± 1.50^{b}	9.00 ± 0.30^{b}	4.35 ± 0.15^{b}	9.00 ± 2.50^{a}	270 ± 0.00^{a}	70 ± 5.00^{a}
WEEK 1	Test	$59.01\pm1.57^{a,A}$	${\bf 39.50} \pm 0.50^{a,A}$	$13.13\pm0.15^{a,A}$	$6.23\pm0.25^{\text{a},A}$	$10.73 \pm 1.25^{a,A}$	$310\pm40.0^{\text{a},\text{BC}}$	$70\pm0.00^{a,B}$
	Control	72.31 ± 3.36^a	32.55 ± 2.95^a	9.90 ± 0.90^a	5.68 ± 0.89^{a}	9.85 ± 5.65^a	335 ± 105.0^a	84 ± 1.40^{a}
WEEK 2	Test	$66.01 \pm 8.84^{a,A}$	$35.15\pm2.05^{a,AB}$	$10.85 \pm 0.75^{a,AB}$	$6.43\pm0.67^{a,AB}$	$12.0\pm3.20^{a,A}$	$333\pm108.5^{a,B}$	$72\pm1.55^{b,B}$
	Control	69.23 ± 2.15^a	32.84 ± 3.95^a	10.36 ± 1.15^a	6.04 ± 0.64^{a}	7.4 ± 2.85^{a}	$423\pm108.0^{\text{a}}$	78 ± 1.40^{b}
WEEK 3	Test	$63.75 \pm 2.55^{b,A}$	$26.23 \pm 3.85^{a,CD}$	$8.15\pm1.35^{a,CD}$	$4.38\pm1.01^{a,B}$	$4.36 \pm 2.50^{a,B}$	$127 \pm 62.50^{a,C}$	$86\pm0.65^{a,A}$
	Control	73.27 ± 2.15^a	39.05 ± 2.35^a	13.83 ± 0.45^a	6.90 ± 1.60^{a}	6.25 ± 0.05^a	416 ± 3.50^{b}	84 ± 0.70^a
WEEK 4	Test	$62.90 \pm 3.84^{b,A}$	$22.50 \pm 1.30^{b,D}$	$6.50\pm0.90^{\text{b},\text{D}}$	$4.36\pm0.15^{a,B}$	$4.33\pm0.11^{\text{b},B}$	$615\pm61.0^{a,A}$	$51\pm2.55^{\text{b},C}$
AVERAGE CONTRO L	Control	69.07 ± 5.9^{A}	30.63 ± 2.8BC	$9.76\pm0.78^{\mathrm{BC}}$	5.31 ± 0.5^{AB}	8.76 ± 3.67^{AB}	$342.83\pm71^{\rm B}$	77.53 ± 2.6^{AB}

 192^{-a-b} Different letters in the same column indicate significant difference (P<0.05) within each week

^{A-B} Different letters in the same column indicate significance difference (P<0.05) across the weeks

195 TABLE 3.3: RESULT OF THE EFFECT OF TOOTHPASTE ON SPERM COUNT

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		SPERM COUNT (X10 ⁶)
	Control	475 ± 125^{a}
WEEK 1	Test	$455\pm5^{b,A}$
	Control	575 ± 25^{a}
WEEK 2	Test	$225\pm225^{b,A}$
	Control	450 ± 150^a
WEEK 3	Test	$125\pm125^{a,A}$
	Control	650 ± 50^{a}
WEEK 4	Test	$250\pm250^{b,A}$
AVERAGE CONTROL	Control	$500 \pm 100^{\mathrm{A}}$

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^{a-b} Different letters in the same column indicate significant difference (P<0.05) within each week

^{A-B} Different letters in the same column indicate significance difference (P<0.05) across the weeks

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203 4.0 DISCUSSION

ALT and AST are general traditional biomarkers used widely for detecting drug induced liver 204 205 injury (Yukuta et al., 2004). In this study, increase or decrease in the levels of these 206 biomarkers is defined by comparing the values obtained from the test animals with the control. The liver enzyme assay showed a gradual increase in the serum levels of AST and 207 ALT with a significant difference in AST (P < 0.05) while there was also a significant 208 209 difference in ALT (P < 0.05) the increase in the level of serum AST and ALT is an indicator 210 of increased activity of the liver possibly due the abnormal presence of sodium fluoride 211 (NaF), Sodium lauryl sulphate (SLS) and other components of the toothpaste that are foreign 212 to the body system. The results also showed that there was a slight significant difference (P< 0.05) in protein and there was a decrease in the protein levels of the test rats as compared to 213

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the control, this might be due to possible negative effect on NaF on the Liver. This decrease 214 215 although inconsistent with the work of (Green, 2005) is consistent with the work of 216 Debensten et al., (1995), Barbar et al. (2006) and Anamika et al, (2012). A more recent study done by Imtithal and Baraa, (2017) indicated that sodium fluoride caused a significant 217 218 decrease in serum protein and albumin concentrations. The general low value of sodium (Na), 219 potassium (K), bicarbonate in the test compared to the control respectively with a significant 220 difference (P < 0.05) on the last week and also when comparing the weeks with the average 221 control might be because of increased secretion of the electrolytes from the body during urine 222 formation. The toothpaste components may cause abnormal inhibition of release of hormones 223 (Anti-Diuretic Hormone) that regulates electrolyte balance. This is because fluoride has been 224 shown to negatively affect the thyroid gland that plays a major role in controlling our body 225 metabolism and internal homeostasis (Bhathnagar et al., 2005), and exposure to it according 226 to Gopalakrishna et al., (2002) can induce the loss of neuronal cell bodies and damage 227 synaptic structures in different regions of the brain. The low level of leukocytes (WBC) 228 recorded on the third and fourth week when compared to the control might be linked to the 229 inflammatory effects of Sodium Fluoride on lymphatic organ, this is in agreement with 230 Maryam et al, (2017). The gradual decrease in PCV, Hb and RBC from week two to week 231 four indicates that NaF has a negative effect on blood when introduced into the system over a 232 long period of time although the difference wasn't significant since Maryam *et al*, (2017) also 233 reported a significant lower blood indices in their experiment. For the sperm count, Results 234 from week 1 to week 4 all had a lesser value of sperm count on the test when compared to the 235 control, with a significant difference (P < 0.05) between the control and the treatment, this 236 significant negative effect of NaF is in agreement with the work of Wang et al., (2006) who 237 reported a deleterious effect of fluoride on the testis which is the site for sperm production, 238 and also agreed with work done by Chinoy and Sequira, (1989) and Arora et al., (2010) who

observed in their experiment that there was a significant decrease in the epididymal sperm
count when sodium fluoride which is a major component of toothpaste was administered to
rats.

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243 5.0 CONCLUSION

244 The results from the study clearly points out that a prolonged ingestion of toothpaste

245 generally affects the function of the liver, kidney and also the semen (for males) negatively

which might lead to renal dysfunction and infertility in men. Based on this efforts should be

- 247 made to prevent the accidental ingestion of toothpaste especially in children.
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