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Ameliorative potential of vitamin E on the impact of dietary fumonisin B_1 on reproductive performance of female rabbits

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Abstract

Fumonisin B_1 (FB₁), a contaminant of agricultural products, particularly maize worldwide is known to be consumed by farm animals and has been documented to cause various physiological responses in animals. A 15-week trial on the ameliorative potential of vitamin E on the negative impacts of FB₁ on reproductive performance of rabbits was conducted. Forty-nine female rabbits aged 16 to 18 weeks weighing 1.65 to 2.0 kg body weight were assigned to seven experimental feeding groups: the control group received a diet without FB₁, three groups were fed diets containing different concentrations of FB₁ at 2.5, 5.0 or 7.5 mg kg⁻¹, and three further groups had diets containing FB₁ and vitamin E i.e., 2.5 mg FB₁ kg⁻¹ + 100 mg vitamin E, 5.0 mg FB₁ kg⁻¹ + 100 mg vitamin E, and 7.5 mg $FB_1 kg^{-1} + 100 mg$ vitamin E. Data obtained on reproductive parameters - gestation length, litter size, kit weight as well as the kit crown-rump length, were analysed using ANOVA. Serum reproductive hormones - luteinizing hormone (LH), follicle stimulating hormone (FSH), prolactin, prostaglanding $F_2\alpha$ (PGF₂ α), and estradiol (E₂) levels in rabbits fed diets containing 7.5 mg FB₁ kg⁻¹ were significantly lower than those for all other treatments. Rabbits fed diets containing $\geq 5 \text{ mg FB}_1 \text{ kg}^{-1}$ had significantly (p < 0.05) longer gestation lengths and lower kit weights compared to the other treatment groups. The litter sizes of rabbits fed FB₁-contaminated diets supplemented with vitamin E were significantly (p < 0.05) higher compared with those on diets not supplemented, including the control. The 21-day *postpartum* weight gain of kits of does fed diets containing $\geq 5.0 \text{ mg FB}_1 \text{ kg}^{-1}$ were significantly (p < 0.05) lower than the weight gain observed in the other treatment groups. Does fed diets supplemented with vitamin E had significantly (p < 0.05) higher milk yield compared with does on not supplemented diets, including the control. This study has shown that vitamin E supplementation of does counteracts the adverse impacts of FB₁ on reproductive hormones, gestation length, kit weight, and milk production in rabbits.

Keywords: rabbit, fumonisin B1, mycotoxin, reproduction, antioxidant

1 Introduction

Rabbit production has potential in many developing countries as a means of supplying cheap, high quality animal protein within the shortest time possible (Oy-awoye *et al.*, 1990). Schiere (2004) stated that growth rate of around 15–20 g day⁻¹ are common in the tropics but it is possible to obtain 30–40 g day⁻¹ if rabbits are

well fed. Since good growth rate and reproductive performance form the basis of high profit margin in livestock production, good nutrition is essential. However, some low-cost rabbit feed constituents, such as maizemilling waste, under humid tropical environmental conditions, may be infected with moulds and consequently may contain mycotoxins. Also, crops contaminated with high concentrations of mycotoxins are often diverted into animal feeds, thereby posing a serious threat to the growth, health and productivity of the animals (Griessler & Encarnação, 2009).

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Fumonisin B₁ (FB₁), a mycotoxin produced by *Fusarium verticillioides* (= *F. moniliforme*) and other *Fusarium* species that grow on cereals, especially maize, has been documented to cause various physiological responses in humans and animals. The effects of dietary FB₁ on reproductive processes in animals have been well documented (Voss *et al.*, 2006; Gbore & Egbunike, 2008; Gbore, 2009a, b; Ewuola & Egbunike, 2010; Gbore *et al.*, 2012).

Considerable research has been directed at finding means to prevent or lessen the toxicity of FB_1 . The use of feed additives is an approach that is considered to be cheaper than degradation of the mycotoxin. Several studies have shown that a variety of adsorbent materials have a high affinity for binding mycotoxins by the formation of stable linkages (Huwig et al., 2001; Galvano et al., 2001; Diaz et al., 2004; Var et al., 2008). Experimental results obtained with some extensively studied adsorbents, such as hydrated sodium calcium aluminosilicate (HSCAS), are quite satisfactory with respect to aflatoxins, but they are not effective in preventing toxic effects of Fusarium mycotoxins, such as fumonisins, trichothecenes or zearalenone (Avantaggiato et al., 2005). In the case of less- and non-absorbable mycotoxins such as these, new strategies have to be applied. Most of the mycotoxins provoke oxygen free radical formation (Balogh et al., 2007; Pál et al., 2009). As a result, the addition of natural or synthetic antioxidants in diets contaminated with these mycotoxins have been reported (Rogers, 2003; Citil et al., 2005; Surai, 2006; Dvorska et al., 2007) to be hypothetically effective in ameliorating the adverse impacts of these mycotoxins on animals due to their superoxide anion scavenging ability.

Dietary concentration of 7.5 mg fumonisin kg⁻¹ feed reportedly impaired reproductive performance in rabbits in 175 day studies (Ewuola & Egbunike, 2010). However, consumption of FB₁ at levels below this threshold may exert suboptimal reproductive performance in breeding female rabbits. The present study was therefore aimed at evaluating the ameliorative potential of vitamin E on the effects of dietary FB₁ below and above the no-observed-adverse-effect-level (NOAEL) of 5 mg kg⁻¹ feed on reproductive performance of female rabbits.

2 Materials and methods

2.1 Animals and experimental site

Forty-nine mixed breeds female rabbits aged 16 to 18 weeks were used in the current study. Rabbits were pro-

cured from a reputable commercial farm in Akure, Ondo State, Nigeria. This study was conducted in the Rabbit Unit of the Teaching and Research Farm of the Federal University of Technology, Akure, Nigeria. The farm is located in the humid rain forest zone of western Nigeria with mean rainfall, relative humidity and temperature of 1,500 mm, 75% and 29 °C, respectively. The raining season is usually from March to November yearly. This study was carried out in accordance with "Guide for the care and use of Laboratory Animals" (NRC, 1996), and approved by the local Institutional Animal Ethics Committee.

2.2 Fumonisin B₁ production and experimental diets

Maize grits cultured with a toxigenic strain of F. verticillioides (MRC 286) and quantified in replicates for FB1 and other common Fusarium mycotoxins as outlined by Gbore et al. (2016) were combined with noncontaminated maize grits to formulate seven diets consisted as follows: a control diet without FB1 contamination, three diets containing different concentrations of FB₁ at 2.5, 5.0 or 7.5 mg FB₁ kg⁻¹ feed, and three further diets containing FB1 and vitamin E at a fixed rate of 100 mg kg⁻¹ contaminated feed (i.e., 2.5 mg FB₁ kg⁻¹ + 100 mg vitamin E, 5.0 mg $FB_1 kg^{-1}$ + 100 mg vitamin E, and 7.5 mg FB₁ kg⁻¹ + 100 mg vitamin E). The level of FB1 in the control diet was below the detection limit of 0.2 mg kg⁻¹ for the mycotoxin. The diets were marked A, B, C, D, E, F, and G, respectively, and provided nearly15 % crude protein, 10 % crude fibre and 2600 kcal of digestible energy kg^{-1} (Table 1).

Table 1: Ingredient and chemical composition (%) of the basal diet.

Ingredient	%
Maize [†]	40.7
Groundnut cake	11.8
Wheat offal	22.6
Palm kernel cake	19.6
Fish meal	2.0
Dicalcium phosphate	2.0
Salt (NaCl)	0.2
DL-Methionine	0.1
L-Lysine	1.06
Minerals/vitamins premix	0.1
Analysed nutrients	
Crude protein (%)	15.46
Crude fiber (%)	9.60
Digestible energy (kcal kg ⁻¹)	2597.96
[†] Varied proportion of <i>Fusarium</i> -conta non-contaminated grains	minated and

2.3 Treatments and experimental layout

The rabbits were weighed at the end of a two week pre-experimental physiological adjustment period and randomly assigned to each of the diets (n = 7 rabbits per treatment) and housed individually in hutches. The animals were fed with their respective experimental diets for 15 weeks.

2.4 Evaluation of reproductive performance of the does

After eight weeks of feeding the respective experimental diets, the rabbits were mated to intact bucks, at the ratio of one buck to two does. At parturition, gestation length, mean litter size, kit weight, and kit crown rump length were determined. The crown rump length of each of the kits was determined using thread calibrated in cm.

2.5 Determination of serum reproductive hormones

Serum level of each of luteinizing hormone (LH), follicle stimulating hormone (FSH), prolactin, prostaglanding $F_2\alpha$ (PGF₂ α), and estradiol (E₂), was determined using appropriate test kits (Endocrines Technologies Inc., Newark, CA). All samples were run in triplicate in a single assay (n=7 per treatment).

2.6 Evaluation of post-partum performance

Rabbits were maintained on the respective diets through gestation till 21 days *post-partum* and the following parameters were determined: litter sizes, kit weights, and mortalities of the offspring at birth, 7, 14 and 21 days *post partum*. Milk yield was calculated using the formula proposed by Lebas *et al.* (1986) i. e., Milk production of does (g) = (Live weight new born at 21 days of age – Live weight of new born)*1.18.

2.7 Statistical analysis

The design used for this experiment is Completely Randomized Design (CRD). Data obtained were subjected to statistical analysis using one-way ANOVA procedure of Statistical Analysis Systems (SAS, 2008). Duncan's multiple range tests of the same software was used to separate all means at 5 % probability level. Results giving *p*-values of < 0.05 were considered significantly different.

3 Results

3.1 Reproductive performance

Dietary FB₁ generally influenced the concentrations of serum hormones examined (Table 2). The serum LH,

FSH, prolactin, $PGF_2\alpha$, and E_2 levels in rabbits fed diets containing 7.5 mg FB₁ kg⁻¹ were significantly lower than those fed on the other treatment diets.

The reproductive performance of the does fed diets containing FB_1 and vitamin E is shown in Table 3. The gestation length of the rabbits was significantly (p < 0.05) influenced by the dietary treatments. Rabbits fed diets containing $\geq 5 \text{ mg FB}_1 \text{ kg}^{-1}$ had significantly longer gestation period compared with does in other treatment groups. The litter sizes of rabbits fed FB₁contaminated diets supplemented with vitamin E were significantly higher (p < 0.05) compared with those on diets not supplemented with vitamin E, including the control. The total and relative weights of the kits of rabbits fed diets containing $\geq 5.0 \text{ mg FB}_1 \text{ kg}^{-1}$ were significantly lower (p < 0.05) than of those in other treatment groups. The crown-rump lengths of kits of rabbits fed diets containing $2.5 \text{ mg FB}_1 \text{ kg}^{-1}$ were significantly (p < 0.05) longer than those of the kits of rabbits in the control group and those on the diet containing 5.0 mg $FB_1 kg^{-1}$.

3.2 Pre-weaning performance

Table 4 shows the pre-weaning performance of kits of does fed different diets. Weight gained by the kits was significantly (p < 0.05) influenced by the dietary treatments. The mean weight of kits of does fed diets containing 7.5 mg FB₁ kg⁻¹ (Diet D) was significantly lower compared to those fed the control diet (Diet A), diets containing 2.5 mg FB₁ kg⁻¹ (Diet B) and \leq 5.0 mg FB_1 kg⁻¹ supplemented with vitamin E (Diets E and F) at day 7 post-partum. At 21-day post-partum, weight gained by kits of does fed diets containing $\geq 5.0 \text{ mg}$ FB_1 kg⁻¹ (Diets C and D) were significantly lower than the weights gained by kits of does in the other treatment groups, with the kits of does fed diets containing 2.5 mg FB_1 kg⁻¹ supplemented with vitamin E having the highest gain. The gain in weights of kits of does fed the control diet, diet containing $2.5 \text{ mg FB}_1 \text{ kg}^{-1}$ and diets supplemented with vitamin E were significantly higher than those of kits of does fed diets containing 5.0 and 7.5 mg FB₁ kg⁻¹ without vitamin E supplementation.

The estimated milk yield of does fed diets containing FB₁ or with vitamin E is shown in Fig. 1. Milk yield of the does was significantly (p < 0.05) influenced by the dietary treatments. Does fed diets supplemented with vitamin E had significantly (p < 0.05) higher milk yield compared with those on diets not supplemented, including the control. The milk yield of the does significantly decreased with increase in the concentrations of FB₁ in the diets.

Parameters $(ng ml^{-1})$	<i>Diet A</i> Control	Diet B 2.5 mg FB ₁	Diet C 5.0 mg FB ₁	Diet D 7.5 mg FB ₁	<i>Diet E</i> 2.5 mg FB ₁ + Vitamin E	<i>Diet F</i> 5.0 mg FB ₁ + Vitamin E	<i>Diet G</i> 7.5 mg FB ₁ + Vitamin E	± SEM
Luteinizing hormone	5.23 ^a	5.20 ^a	5.20 ^a	4.53 ^b	5.00 ^{<i>a</i>}	5.27 ^a	5.00 ^{<i>a</i>}	0.15
Follicle stimulating hormone	17.37 ^a	17.80 ^a	17.00 ^a	14.20 ^b	17.00 ^a	17.23 ^a	17.93 ^a	0.69
Prolactin	2.03 ^a	2.00 ^a	1.90 ^{<i>a</i>}	1.53 ^b	2.10 ^{<i>a</i>}	2.03 ^a	2.07 ^a	0.15
Prostaglandin $F_2 \alpha$	5.77 ^a	5.70 ^{<i>a</i>}	5.83 ^a	4.33 ^b	5.93 ^a	5.93 ^a	5.77 ^a	0.28
Estradiol	9.11 ^{<i>a</i>}	9.75 ^a	9.38 ^{<i>a</i>}	6.52 ^b	9.47 ^a	9.42 ^{<i>a</i>}	9.33 ^a	0.66
^{<i>a,b</i>} : Means on the same row with different superscripts differ significantly ($p < 0.05$).								

 Table 2: Serum hormones of female rabbits fed diets containing FB1 with or without vitamin E.

Table 3: Reproductive performance of does fed diets containing FB_1 with or without vitamin E (Mean \pm SE).

Parameters	Diet A Control	Diet B 2.5 mg FB ₁	Diet C 5.0 mg FB ₁	Diet D 7.5 mg FB ₁	Diet E 2.5 mg FB ₁ + Vitamin E	Diet F 5.0 mg FB ₁ + Vitamin E	Diet G 7.5 mg FB ₁ + Vitamin E
Gestation length (d)	30.33±0.58 ^c	30.00±0.00 ^c	32.00±1.00 ^b	33.33±1.53 ^a	29.67±0.58 ^c	30.00±0.00 ^c	30.00±0.00 ^c
Litter size	4.67±1.15 ^{ab}	3.67±1.53 ab	3.67±0.58 ^{ab}	3.00 ± 1.00^{b}	5.33±1.15 ^a	5.33±1.15 a	5.33±0.58 ^a
Litter weight (g)	224.80±57.43 ^{ab}	173.30±80.68 ^b	116.67±16.04 ^c	119.97 ± 36.64 ^c	284.60 ± 61.35^{a}	247.40±57.16 ^{ab}	235.57±11.05 ab
Kit weight (g)	45.83±4.01 ab	48.13±2.11 ab	46.23±5.40 ^{ab}	40.30±1.65 ^b	53.77±6.76 ^a	46.30±1.76 ^{ab}	53.83±4.27 ^a
Kit crown rump length (cm)	12.00±1.00 ^b	13.33±0.76 ^a	12.00±0.50 ^b	10.67±0.58 ^c	10.67±0.58°	10.33±0.58 ^c	11.33±0.58 ^{bc}
a,b,c. Means on the same row with different superscripts differ significantly ($p < 0.05$)							

Table 4: *Pre-weaning performance of kits of does fed diets containing* FB_1 *with or without vitamin E (Mean* \pm *SE).*

Parameters	Diet A Control	<i>Diet B</i> 2.5 mg FB ₁	Diet C 5.0 mg FB ₁	Diet D 7.5 mg FB ₁	<i>Diet E</i> 2.5 mg FB ₁ + Vitamin E	<i>Diet F</i> 5.0 mg FB ₁ + Vitamin E	<i>Diet G</i> 7.5 mg FB ₁ + Vitamin E
Weight at birth (g)	45.83±4.01 ab	48.13±2.11 ab	46.23±5.40 ^{ab}	40.30±1.65 ^b	53.77±6.76 ^a	46.30±1.76 ^{ab}	53.83±4.27 ^a
Weight at day 7 (g)	83.53±6.76 ^{ab}	80.33±0.58 ^{ab}	77.80±7.12 ^{bc}	68.20±6.18 ^c	90.60±8.93 a	88.57 ± 5.56^{ab}	78.83±2.76 ^{bc}
Weight at day 14 (g)	118.10±2.79 ^{ab}	$117.87 {\pm} 2.58^{ab}$	115.90±6.50 ^{ab}	99.40±11.46 ^c	127.27±5.99 ^a	123.67±4.56 ^a	120.40±7.24 bc
Weight at day 21 (g)	139.33±11.41 ^{bc}	135.00±5.00°	116.27 ± 7.11^{d}	114.07 ± 8.93^{d}	155.30±3.12 ^a	142.43±3.82 ^{ab}	140.07±5.05 ab
Change in weight (g)	93.50±4.02 ^b	86.87±2.11 ^c	70.04 ± 4.50^{d}	73.77 ± 1.50^{d}	101.53±6.70 ^a	96.13±1.75 ^b	86.24±3.22 ^c
Mortality at Birth	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	$0.00 {\pm} 0.00$	0.00 ± 0.00	$0.00 {\pm} 0.00$	0.00 ± 0.00
Mortality at day 7	1.00 ± 1.00	1.33 ± 0.58	0.33±0.58	1.33 ± 0.58	0.33 ± 0.58	1.00 ± 0.00	0.33 ± 0.58
Mortality at day 14	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.33 ± 0.58	0.00 ± 0.00	$0.00 {\pm} 0.00$	0.00 ± 0.00
Mortality at day 21	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	$0.00 {\pm} 0.00$	0.00 ± 0.00	$0.00 {\pm} 0.00$	0.00 ± 0.00
a,b,c,d: Means on the same row with different superscripts differ significantly ($p < 0.05$).							

4 Discussion

Although, previous studies (Voss *et al.*, 1996; LaBorde *et al.*, 1997; Collins *et al.*, 1998a,b) provided no evidence that FB₁ is a reproductive toxicant, more recent observations and experimental findings have however shown that FB₁ is a possible risk factor for birth defects (Merrill *et al.*, 2001; Marasas *et al.*, 2004; Voss *et al.*, 2006) and impaired reproductive capacity in animals (Gbore & Egbunike, 2008; Gbore, 2009a, b; Ewuola & Egbunike, 2010; Gbore *et al.*, 2012; Cortinovis *et al.*, 2014; Albonico *et al.*, 2016). In the present study, results show that > 5 mg dietary $FB_1 kg^{-1}$ feed had observable adverse effect on gestation length in rabbits. However, the inclusion of vitamin E at 100 mg kg⁻¹ diet countered this effect by reducing the gestation length. This could be attributed to the fact that vitamin E plays an important role in reproductive performance in animals due to its anti-oxidative properties which enabled its participation in metabolism of all cells as reported by Gutteridge & Halliwell (1994). Seemingly elongated gestation lengths was observed in rats fed *F. monili-forme* culture material containing 10 ppm FB₁ from two



Concentrations of FB_1 or with vitamin E (per kg of diet)

Fig. 1: Mean milk yield of does.

weeks before mating by Voss *et al.* (1996). In a previous study, Gbore *et al.* (2012) observed significantly elongated gestation length from 21.60 ± 0.53 days in rats fed diets containing $\geq 10.0 \text{ mg FB}_1 \text{ kg}^{-1}$ to 23.33 ± 0.51 days. These reports and the results from this study, further lead credibility to the fact that FB₁ potentially affects reproductive development in rabbits. The result of this study on gestation length shows that FB₁ at low concentration of 2.5 mg kg^{-1} had no detrimental effect on the performance of rabbits but at higher concentrations of 5.0 and 7.5 mg kg⁻¹ the detrimental effects were observable.

Gbore *et al.* (2012) reported depressed serum gonadotropins levels in rats fed diets containing $\geq 10 \text{ mg}$ FB₁ kg⁻¹. The authors postulated that this might be due to an increase in gonadal steroid inhibition or suppression of the hypothalamus and/or pituitary gland with resultant decline in serum gonadotropins levels. Adverse effects of mycotoxins on sexual and reproductive developments have been reported. Dietary zearalenone levels as low as 0.05–0.06 mg kg⁻¹ DM have been shown to increase the number of ovarian follicles and to decrease the serum concentration of the gonadotropic hormone (FSH) in female piglets (Döll *et al.*, 2003), thus potentially affecting their sexual development. Also, dietary FB₁ was reported to delay attainment of sexual maturity in growing pigs (Gbore, 2009a) and rabbits (Ewuola & Egbunike, 2010). Fertility disturbances and other reproductive pathologies, particularly suppressive effect on testosterone secretion in mice, was reported (Yang *et al.*, 2007 a,b) following ingestion of cereals contaminated with *Fusarium* fungi. Also, reduced progesterone synthesis due to inhibition of the follicle stimulating hormone secretion (FSH) by *Fusarium* mycotoxins in cultured granulosa cells (GCs) from porcine ovaries was reported by Tiemann *et al.* (2003).

Improved litter size in rabbits fed diets supplemented with vitamin E over other treatment groups in this study may be attributed to the role vitamin E plays in reproduction. In the present study, it is evident that the dose-dependent significant decline in litter weight of does fed diets B, C, and D was countered by vitamin E supplementation. In a study, LaBorde *et al.* (1997) reported a reduced foetal weight from rabbits dosed daily by gavage on gestation days (GD) 3–19 with purified FB₁ at 0.5–1 mg kg⁻¹ day⁻¹. The authors ascribed this weight reduction to maternal toxicity, rather than any developmental toxicity produced by FB₁. However, FB₁ dose-responsive significant decrease in foetal weight observed in this study correlates with reports of other studies. Pregnant rats dosed by gavage on GD 8– 12 with a semi-purified extract of culture material containing FB₁ with a purity of 80% resulted in lower foetal weight at dose of 60 mg kg⁻¹ (Lebepe-Mazur *et al.*, 1995). Similarly, decreased body weight of live foetuses obtained from pregnant Syrian hamsters dosed with FB₁ in a dose-dependent manner was reported in a study by Penner *et al.* (1998). Voss *et al.* (1996) reported lower litter weights from rats fed *F. moniliforme* culture material providing 1–55 ppm FB₁ from two weeks before mating compared to the control group.

The significantly reduced litter weights at concentration of 7.5 mg FB₁ kg⁻¹ in this study corroborates the report of Voss et al. (1996) that FB1 at high concentrations of 10 and 55 mg kg⁻¹ significantly reduced the weight of dams fed diets contaminated with F. verticillioides. It is also similar to the report of Lebepe-Mazur et al. (1995) that FB_1 at high dose significantly suppressed growth in rats. In the present study, vitamin E countered this effect by increasing the litter weight. The significant positive impact of vitamin E could be a result of the crucial role it plays in the growth of animals. It has been reported that FB_1 is not transferred through the placenta or into the milk in several animal species (Scott et al., 1994; Becker et al., 1995; Voss et al., 1996; LaBorde et al., 1997; Collins et al., 1998b), nor are its metabolites found in animal products such as milk, meat and eggs (Jonker & van Egmond, 1999). Therefore, the resultant depressed growth rate of litters of does fed diets containing $\geq 2.5 \text{ mg FB}_1 \text{ kg}^{-1}$ without vitamin E supplementation may be attributed to the reduced milk yields in does fed diets B, C, and D (Fig. 1).

Exposure to Fusarium mycotoxins has been linked to reproductive disorders in pigs (Cortinovis et al., 2014; Gbore & Egbunike, 2008; Gbore, 2009a,b), mice (Yang et al., 2007a,b), and rabbits (Ewuola & Egbunike, 2010). In a study to determine the potential reproductive effect of FB₁ on granulosa cell (GC) proliferation and steroid production in swine, Cortinovis et al. (2014) reported that this mycotoxin alone or with other Fusarium mycotoxins, including deoxynivalenol (DON) and zearalenone, influenced porcine GC proliferation and steroid production, thereby demonstrating their potential reproductive effects on swine. GC is reported to be crucial in the process of normal folliculogenesis and oocyte growth and development as they provide essential nutrients to the oocyte and establish a link between the oocyte and the surrounding ovarian tissue in which the follicle is embedded (Petro et al., 2012). Also, GC is responsible for ovarian steroidogenesis that can be altered at any level leading to changes in the rate of hormone production and concentration (Ndossi et al., 2012; Petro et al., 2012). Progesterone is an essential regulator of the reproductive events and plays key roles in ovulation, zygote implantation, and subsequent maintenance of pregnancy (Graham & Clarke, 1997). The influence of FB₁ on porcine GC function as noted by Cortinovis *et al.* (2014) could be one mechanism whereby this mycotoxin may impair reproductive activity in animals, particularly inhibiting effect on synthesis and secretion of gonadotropins. In rats, FB₁ concentrations at $\geq 10 \text{ mg kg}^{-1}$ diet were found to significantly reduce serum gonadotropin levels without inducing histopathological changes in the ovaries (Gbore *et al.*, 2012).

There exists a high correlation between the milk production by the doe and the growth of the kits because rabbit kits do not show significant feed intake before the age of 18-19 days (Fortun-Lamothe & Gidenne, 2000). Although Pascual et al. (1996) reported increase in milk yield with increase in litter sizes, the significantly decreased milk yield with increase in the concentrations of FB₁ in does fed diets supplemented with vitamin E, which had similar litter size, could be attributed to the mycotoxin in the diets. FB1 bears a remarkable structural resemblance to sphinganine and sphingosine and the mycotoxin inhibits sphingosine metabolism in tissues, leading to an accumulation of sphingoid bases, which are intermediates in sphingolipid biosynthesis (Wang et al., 1991). Sphingolipids are critical components for cell survival and homeostasis (Hirabayashi, 2012). High levels of sphingolipids have been reported in brains (Assi et al., 2013) and its metabolism has a key neuropathological impact (Mencarelli & Martinez-Martinez, 2013). The disruption of sphingosine metabolism and alterations in the amounts of any of these intermediates by FB₁, which could potentially alter neuronal function (Fonteh et al., 2006) and result in a variety of biological effects (Penner et al., 1998), may be responsible for the significant decline in serum gonadotropins and consequently reduced milk yield of does, with increased dietary FB₁ as observed in this study.

5 Conclusion

Economic loss in livestock production due to mycotoxins in animal feeds is a major problem in the tropics, including Nigeria. This study revealed that reproductive and pre-weaning performances of rabbits exposed to diets containing FB₁ could be impaired at concentrations above 5.0 mg kg⁻¹ of feed. Inclusion of vitamin E at 100 mg kg⁻¹ of feed in FB₁-contaminated feeds is recommended as a nutritional strategy to alleviate the reproductive-depressing effect of the mycotoxin in female rabbits, especially at concentrations of ≥ 5.0 mg FB₁ kg⁻¹.

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