

## HAEMATOLOGICAL PARAMETERS OF WEST AFRICAN DWARF SHEEP AND GOAT FED A BASAL DIET OF CASSAVA PEEL SUPPLEMENTED WITH FRESH *GLIRICIDIA SEPIUM* LEAVES

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**Abstract:** This study was conducted to investigate the effects of feeding fresh *Gliricidia sepium* leaves as supplement to basal diet of dried cassava peel on the haematological parameters of West African Dwarf sheep and goats in a completely randomized design experiment. Twelve each of WAD sheep and goat were randomly divided into three groups of four animals per group for both sheep and goats. The three groups were given different diets of (T<sub>1</sub>: 100% cassava peel, T<sub>2</sub>; 70% cassava peel + 30% *Gliricidia* and T<sub>3</sub>; 50% cassava peel + 50% *Gliricidia*) for a period of 7 weeks with an initial 2 weeks of acclimatization. Haematological indices of these animals were investigated; before and after experiment. Parameters monitored included; PCV, WBC, RBC, Hb, Neutrophil and lymphocytes. The results showed that cassava peel was higher in DM than *Gliricidia* but *Gliricidia sepium* had higher values of CP, EE and total Ash than cassava peel. Animals fed 100% cassava peel (T<sub>1</sub>) had better PCV, WBC and Hb values followed by T<sub>3</sub> and T<sub>2</sub> in goat. Similarly, PCV and Hb values in sheep were higher under T<sub>1</sub> while WBC, RBC and lymphocytes had better values in T<sub>3</sub>.

Both species of animals (sheep and goat) can be fed dried cassava peel supplemented with fresh leaves of *Gliricidia sepium* at 50% inclusion level and perform well during the dry season. Supplementation with *Gliricidia sepium* will cause no harm on the performance of sheep and goat in term of haematological parameters, most especially those investigated in this study.

**Keywords:** Cassava peel, WAD, Haematological parameters, Basal diet, *Gliricidia*.

### Introduction

Livestock productivity in the tropics has suffered major setback due to inadequate quantity and quality feeds for the animals especially during dry season (Peters, 1998). In wet season, forage is relatively available and animals (small ruminants) may easily gain weight and remain thrifty (Babayemi, *et al.*, 2003). Akinlade, *et al.*, (2005) reported that a major problem facing small ruminant animal producer is how to feed the animals adequately in all year round. Small ruminant production in dry season could be improved upon; by cultivating forage plants with high leaf yield (Fasae; *et al.*, 2009). These crops when harvested, could be

dried or stored as food for ruminants during the scarce period when these animals lose a considerably percentage of their body weight as a result of shortage of good quality forage.

In meeting the energy, protein, crude fibre and mineral requirements of small ruminant animals during dry season period, Adegbola and Asaolu (1985) identified cassava peel as an important by product. Browse plant with high nutritive value have been successfully fed to ruminant animals in alley farming system (Fasae and Alokun, 2006). Studies have shown that multipurpose trees can be used as cheap protein supplement which can improve voluntary intake, digestibility and general performance of animals fed low quality feeds. *Gliricidia sepium* has been described as suitable feed for ruminants which can be consumed in large quantities without deleterious effects on animal performance (Bawala, *et al.*, 2006). It contains nutrient levels which are superior to minimum critical levels required for ruminants, and is comparable to other shrubs considered as having a right fodder quality (Smith and Van Houtert, 1981). The nutritive value of *Gliricidia sepium* as animal feed had been studied by Carew (1982), Mba *et al.*, (1981), Onwuka (1980) among others. These studies indicated that *Gliricidia sepium* contains between 3.2 to 4.2% N and can also be used as a source of supplemental energy in animal feeding.

Blood is a specialized bodily fluid in animals that delivers necessary substances such as nutrients and oxygen to the cells and transport metabolic waste products away from those same cells. In vertebrates, it is composed of blood cells suspended in a liquid called blood plasma. Plasma, which constitute 55% of blood fluid, is mostly water (92% by volume) (Franklin institute, 2009) and contains dissolved proteins, glucose, mineral ions, hormones, carbon-dioxide (plasma being the main medium for excretory products transportation) and blood cells themselves. Blood performs many important functions within the body including: supply of oxygen to tissues (bound to haemoglobin, which is carried in red cells, supply of nutrients such as glucose, amino acids and fatty acids (dissolved in the blood or bound to plasma proteins e.g. blood lipids), removal of waste such as carbon dioxide, urea and lactic acid, hydraulic functions and Regulation of core body temperature (Franklin institute, 2009).

## **Materials and Methods**

### **Experimental Site**

The experiment was carried out at the small ruminant unit of the Ladoké Akintola University of Technology Teaching and Research Farm, Ogbomoso. The area is located at 8° 10' North latitude and 4° 10' East longitudes with annual rainfall of 1270 to 2030mm, which occurs in 7–10 months with a peak between July and September of the year. The temperature

of the area ranges between 33<sup>0</sup>C to 38<sup>0</sup>C, with humidity of about 79% all year round except in January when the dry wind blows from North (Olaniyi, 2006).

### **Experimental Animals and their Management**

Twelve West African Dwarf sheep and goats of about one year old weighing 10 – 12kg for sheep, 6 – 8kg for goat were purchased from local markets. They were randomly divided into three treatments of four animals per treatment (for sheep and goat). The animals on arrival were confined in the quarantine pen which had been previously washed and disinfected using Iodophor solution. The animals were treated against internal and external parasites using Ivomec at 1ml to 10kg body weight. They were also given long acting antibiotic at 1ml to 10kg body weight. They were then transferred into individual pens having the floor covered with wood shaving as bedding materials.

### **Experimental Diet**

Cassava peels were collected from a local “garri” processing plant and sundried on a clean concrete floor for 5 days. *Gliricidia sepium* was harvested from existing plantation at the Teaching and Research farm. The leaves were pooled together and air dried to a constant moisture level and thereafter bagged for later use. The animals were allotted experiment as follows; T<sub>1</sub> was given 100% cassava peel, T<sub>2</sub> 70% cassava peel + 30% *Gliricidia sepium* and T<sub>3</sub> 50% cassava peel + 50% *Gliricidia sepium*. Each animal within a group received 5% of its body weight daily of an assigned diet for 7 weeks and water was given ad libitum.

### **Chemical Analysis**

The feed samples were analysed for their proximate composition to determine their Dry matter (DM), Crude protein (CP) Crude fat, Crude fibre (CF) and Total Ash according to AOAC (1995).

### **Data Collection**

Blood sample was collected from each animal twice; on the 1st day of experiment and the last day of experiment through the jugular vein puncture using sterilized needle and syringe and emptied into collecting tube containing Ethylene-diamine-tetracetic acid (EDTA anti-coagulant). These blood samples were taken to laboratory where the following haematological parameters were determined; Red Blood Cell Count (RBC), White Blood Cell count (WBC), Packed Cell Volume (PCV), Hemoglobin Concentration (Hb), Neutrophil and Lymphocytes.

### Statistical Analysis

Data obtained were subjected to analysis of variance (ANOVA) using the general linear model procedure of SAS 2000. The means were separated using Duncan multiple range test of the same statistical packages.

### Results and Discussions

The chemical composition of cassava peel and *Gliricidia sepium* as shown in table 1 indicates higher Dry matter in cassava peel than what was obtained in *Gliricidia sepium* leaves. These values were higher than (66.25%) reported by Adegbola and Asaolu (1985) for cassava peel and (34.5%) reported by Onwuka (1985) for *Gliricidia sepium*. The crude protein content of cassava peel and *Gliricidia sepium* leaves were lower in value than (5.94%) and (20.69%) respectively, obtained by Onwuka (1985) when he studied *Gliricidia sepium* as dry season feed for goat production in Nigeria but cassava peel value was similar to what was reported by Adegbola and Asaolu (1985). Crude fat in cassava peel and *Gliricidia sepium* leaves were higher than values reported by Onwuka (1985) but the total Ash values were lower to results of Onwuka (1985) and Adegbola and Asaolu (1985). These variation or different values could be attributed to different environment where these feed stuffs were gotten, seasonal variation or time of the year (dry or raining season), age of the plant and varieties.

The experimental diet had significant effect on the haematological parameters of WAD sheep as shown in table 3. Though the WBC, RBC, Hb and Neutrophil values were higher in T<sub>2</sub> and T<sub>3</sub> before experimentation (Table 2) but higher values were obtained in T<sub>1</sub>, T<sub>2</sub> and T<sub>3</sub> for lymphocytes at the end of experiment (Table 3) compared to what was obtained in table 2. Despite these significant effects of treatments, T<sub>1</sub> still retained its value unchanged which showed that if available in abundant during dry season, dried cassava peel (T<sub>1</sub>) can be fed to sheep. The PCV value obtained from T<sub>1</sub>, T<sub>2</sub> and T<sub>3</sub> (Table 3) despite the effects of experimental diet still fell within the normal range for sheep as reported by (Reece and Swenson, 2004). The lower values recorded for WBC and RBC for T<sub>1</sub> (Table 3) could be attributed to low level of protein (CP) in the feed (Table 1). The Hb value ranged from (9.83 – 13.00g/dl) and fell within the normal value reported by Adaramola, *et al.*, (2005). The neutrophil values that were statistically similar ( $p > 0.05$ ) in all treatments were higher than the values (28.25 and 35) reported by Anaeto *et al.*, (2010) who studied the effect of cassava leaf silage and cassava peel diet on WAD sheep during dry season. Meanwhile the

lymphocytes values recorded in this study (Table 3) were lower to the value obtained by Anaeto *et al.*, (2010).

Goat showed positive response to experimental diet in lymphocytes with higher value recorded in T<sub>2</sub> than T<sub>1</sub> and T<sub>3</sub> (Table 5) which were also higher than what were obtained before experimental diet was given (Table 4). The treatment also had positive effect on T<sub>1</sub>forHb and T<sub>3</sub>for WBC than what were obtained in table 4. The PCV range of this study were lower to 32 – 45% reported by Banerjee (2005) but in agreement with what was reported by Adaramola *et al.*, (2005) and higher than 28.30 – 28.80% reported by Ukanwoko and Ironkwe (2012) when they studied the growth performance and haematological values of West African Dwarf (WAD) goats fed *Leucaena*, *Gliricidia* and cassava leaf meal – cassava peel based diets. The PCV value in T<sub>1</sub> (Table 5) which was significantly ( $p < 0.05$ ) different from others counts that cassava peel had ability to sustain goat without much effect on the total blood volume. The WBC ( $10^3/ml$ ) values in T<sub>1</sub> and T<sub>3</sub> which were significantly higher ( $p < 0.05$ ) than T<sub>2</sub> (Table 5) were lower to what was reported by Ukanwoko and Ironkwe (2012). The values recorded in this study for Hb in T<sub>1</sub>, T<sub>2</sub> and T<sub>3</sub> (Table 5) were higher than what was reported by Ukanwoko and Ironkwe (2012) and compared favourably with the range of 7 – 15(g/dl) reported for WAD goats by Adaramola *et al.*, (2005). The Neutrophil, though not significantly different ( $p > 0.05$ ) in all treatments as well as lymphocytes, their values fell within the normal physiological values for WAD goats (neutrophil = 17- 52% and Lymphocytes 47 – 82%) (Daramola *et al.*, 2005) as reported by Ifut *et al.*, (2010).

**Table 1: Proximate Composition of cassava peel and *Gliricidia* leaves**

PARAMETERS	CASSAVA PEEL	<i>GLIRICIDIA SEPIUM</i>
Dry matter (DM) %	88.68	70.84
Crude protein (CP)%	5.35	19.68
Crude fat %	5.11	10.93
Crude fibre %	3.35	2.77
Total Ash	4.77	7.51

**Table 2: Haematological parameters of WAD sheep before experimentation**

Parameter	T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>	SEM
PCV (%)	41.33	37.33	43.00	2.4
WBC(10 <sup>3</sup> /ml)	10533.33	9300.00	10900.00	937.0
RBC(10 <sup>6</sup> /ml)	13600000.00	13866666.67	13600000.00	1288122.3
Hb(g/dl)	13.17	11.93	13.40	0.8
Neutrophil (%)	42.33 <sup>b</sup>	42.67 <sup>b</sup>	58.33 <sup>a</sup>	2.6
Lymphocytes (%)	59.00 <sup>a</sup>	57.33 <sup>a</sup>	41.67 <sup>b</sup>	2.8

<sup>ab</sup>, means within the same row with different superscripts are significantly (P < 0.05) different.

**Where:**

PCV = Packed cell volume, WBC = White blood cell, RBC = Red blood cell, Hb = Haemoglobin T<sub>1</sub> = Treatment 1, T<sub>2</sub> = Treatment 2, T<sub>3</sub> = Treatment 3, SEM = Standard error of mean, WAD = West African Dwarf.

**Table 3: Haematological parameters of WAD sheep after experimentation**

Parameter	T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>	SEM
PCV (%)	41.33 <sup>a</sup>	28.00 <sup>c</sup>	33.33 <sup>b</sup>	1.8
WBC(10 <sup>3</sup> /ml)	6000.00 <sup>b</sup>	6533.33 <sup>b</sup>	8933.33 <sup>a</sup>	449.2
RBC(10 <sup>6</sup> /ml)	5413333.33 <sup>b</sup>	7200000.00 <sup>ab</sup>	8533333.33 <sup>a</sup>	853715.3
Hb(g/dl)	13.00 <sup>a</sup>	9.83 <sup>b</sup>	12.07 <sup>a</sup>	0.4
Neutrophil (%)	39.33	42.00	38.00	2.9
Lymphocytes (%)	60.67	58.00	62.00	2.9

<sup>ab</sup>, means within the same row with different superscripts are significantly (P < 0.05) different.

**Where:**

PCV = Packed cell volume, WBC = White blood cell, RBC = Red blood cell, Hb = Haemoglobin T<sub>1</sub> = Treatment 1, T<sub>2</sub> = Treatment 2, T<sub>3</sub> = Treatment 3, SEM = Standard error of mean, WAD = West African Dwarf.

**Table 4: Haematological parameters of WAD goat before experimentation**

Parameter	T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>	SEM
PCV (%)	40.00 <sup>b</sup>	58.00 <sup>a</sup>	53.00 <sup>a</sup>	2.0
WBC(10 <sup>3</sup> /ml)	17800.00 <sup>a</sup>	17500.00 <sup>a</sup>	13133.3 <sup>b</sup>	1283.9
RBC(10 <sup>6</sup> /ml)	12533333.33 <sup>b</sup>	17333333.33 <sup>a</sup>	11226666.67 <sup>b</sup>	1619712
Hb(g/dl)	12.67 <sup>b</sup>	17.33 <sup>a</sup>	16.33 <sup>a</sup>	0.6
Neutrophil (%)	34.00	39.33	40.33	6.1
Lymphocytes (%)	66.00	60.67	59.00	6.0

<sup>ab</sup>, means within the same row with different superscripts are significantly ( $P < 0.05$ ) different.

**Where:**

PCV = Packed cell volume, WBC = White blood cell, RBC = Red blood cell, Hb = Haemoglobin T<sub>1</sub> = Treatment 1, T<sub>2</sub> = Treatment 2, T<sub>3</sub> = Treatment 3, SEM = Standard error of mean, WAD = West African Dwarf.

**Table 5: Haematological parameters of WAD goat after experimentation**

Parameter	T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>	SEM
PCV (%)	38.67 <sup>a</sup>	30.33 <sup>b</sup>	32.33 <sup>b</sup>	1.7
WBC(10 <sup>3</sup> /ml)	15066.67 <sup>a</sup>	10733.33 <sup>b</sup>	14333.33 <sup>b</sup>	1230.1
RBC(10 <sup>6</sup> /ml)	10666666.67 <sup>a</sup>	7200000 <sup>b</sup>	7200000 <sup>b</sup>	55511
Hb(g/dl)	13.27 <sup>a</sup>	10.67 <sup>b</sup>	11.07 <sup>ab</sup>	0.7
Neutrophil (%)	33.33	26.67	34.00	2.7
Lymphocytes (%)	66.67	73.33	66.00	2.7

<sup>ab</sup>, means within the same row with different superscripts are significantly ( $P < 0.05$ ) different.

**Where:**

PCV = Packed cell volume, WBC = White blood cell, RBC = Red blood cell, Hb = Haemoglobin T<sub>1</sub> = Treatment 1, T<sub>2</sub> = Treatment 2, T<sub>3</sub> = Treatment 3, SEM = Standard error of mean, WAD = West African Dwarf.

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