

Hematological profile of East African Short-Horn Zebu calves from birth to 51 weeks of age

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Abstract

This paper is the first attempt to accurately describe the hematological parameters for any African breed of cattle, by capturing the changes in these parameters over the first 12 months of an animal's life using a population-based sample of calves reared under field conditions and natural disease challenge. Using a longitudinal study design, a stratified clustered random sample of newborn calves was recruited into the IDEAL study and monitored at 5-weekly intervals until 51 weeks of age. The blood cell

analysis performed at each visit included: packed cell volume; red cell count; red cell distribution width; mean corpuscular volume; mean corpuscular hemoglobin concentration; hemoglobin concentration; white cell count; absolute lymphocyte, eosinophil, monocyte, and neutrophil counts; platelet count; mean platelet volume; and total serum protein. The most significant age-related change in the red cell parameters was a rise in red cell count and hemoglobin concentration during the neonatal period. This is in contrast to what is reported for other ruminants, including European cattle breeds where the neonatal period is marked by a fall in the red cell parameters. There is a need to establish breed-specific reference ranges for blood parameters for indigenous cattle breeds. The possible role of the postnatal rise in the red cell parameters in the adaptability to environmental constraints and innate disease resistance warrants further research into the dynamics of blood cell parameters of these breeds.

Introduction

The sedentary mixed crop-livestock smallholding system encompasses >50% of poor people resident in East Africa. These resource-poor subsistence farmers own only a few animals, yet these animals are one of their principal capital assets. Subsistence farmers rely on livestock for food security, financial income and social welfare. Indigenous breeds are regionally very important and constitute up to 77% of the total Kenyan cattle population (Rege et al. 2001). The small East African Short-horn Zebu, in particular, is the most common breed in eastern and parts of south-central Africa (Mwacharo et al. 2006). Indigenous breeds are preferred by farmers for their adaptability in terms of disease resistance, heat tolerance, and water and food requirements (Mwacharo and Drucker 2005). Cattle are used for animal traction, milk and meat, and the dung for fertilizer (Rege et al. 2001).

The tropical climate in western Kenya is conducive for the survival of many infectious pathogens and vectors. The most economically important diseases of livestock in sub-Saharan Africa are tick-borne diseases (East Coast fever (ECF), heartwater, anaplasmosis and babesiosis), trypanosomosis (Uilenberg 1995; Minjauw and McLeod 2003; Maudlin 2006) and helminthosis. Small-holder farmers are particularly vulnerable to the economic impact of infectious diseases of livestock. Generally tick-borne infections do not affect trade in livestock, but they are a significant cause of production losses (Perry and Young 1995). Losses include lowered production rates, mortalities, decreased reproduction rates and costs of treatment and control measures. These diseases also indirectly constrain livestock production through limiting the use of the highly susceptible improved breeds of livestock that are used in other countries to improve livestock productivity (Perry and Young 1995).

Baseline values of the parameters that can be used to measure an animal's health status, including hematological parameters, are lacking for indigenous cattle breeds. Most published reference ranges for these parameters for cattle were compiled for European breeds (Jain 1993, Knowles et al. 2000; Mohri et al. 2007). Studies have indicated that baseline values for hematological parameters differ between the various cattle breeds in the tropics (Oduye and Okunaiya 1971). The aim of this study was to investigate the hematological profile of East African Short-horn Zebu cattle living in tropical field conditions in Western Kenya, in particular the age-related changes in the blood parameters from neonate to 51 weeks. This paper is the first attempt to accurately describe the hematological parameters for any African cattle breed capturing the changes in these parameters over the first 12 months of an animal's life using a population based sample of calves reared in their natural environment and disease challenges.

Materials and Methods

The study was carried out in Busia district, Western Kenya (Fig. 1) and a full description of the design is given by Bronsvort et al. (2012) (in press). This site was considered to

represent of smallholder livestock farming systems in East Africa. The climate is tropical and the main land use in the area is mixed crop/livestock-farming. The main crops cultivated include maize, sugarcane, cotton, pigeon-peas and sisal. The main cattle breed kept is the indigenous Zebu, of which the majority are East African Short-horn Zebus. These Short-horn Zebus are a relatively small breed but are considered to be well adapted to the local environmental constraints (Fig. 2). The calves recruited into the study remained with their herd of origin, and were exposed to the natural pathogen challenge in the field. For the duration of the study no interventions in terms of prevention or treatment (except on welfare grounds in which case calves were censored) against infection were practiced.

A stratified 2-stage cluster sampling strategy was used. Stratification was by agro-ecological zone (AEZ) (Jaetzold and Schmidt 1983) with 1st stage cluster (sub-location) selected using stratified-random sampling with replacement and 2nd stage cluster (calf) selected using ordinary random sampling without replacement. A total of 548 calves from sedentary mixed crop-livestock smallholding farms were recruited between October 2007 and September 2009. Calves were recruited within 7 days of birth and monitored routinely at 5-weekly intervals until 51 weeks of age. At each visit a complete clinical examination of the calf was performed and relevant samples collected. These samples included thin and thick peripheral smears of blood collected from the marginal ear vein, jugular blood in 10 ml plain vacutainer tube for the separation of serum, and jugular blood for hematology in 5 ml ethylene-diamine-tetra-acetic acid (EDTA) plastic tubes.

Initial sample processing and analysis occurred at the local field laboratory, Busia, Kenya. Packed cell volume (PCV) was measured manually using a Hawksley microhematocrit reader (Jain 1993). Automated blood cell analysis was performed using the pocH-100iV Diff (Sysmex© Europe GMBH). The hematological parameters investigated included: red cell count (RCC), hemoglobin concentration (HGB), mean corpuscular volume (MCV), mean corpuscular hemoglobin concentration (MCHC), red cell distribution width (RDW), total

white cell count (WCC), platelet count (Plt) and mean platelet volume (MPV). Total serum protein (TSP) was measured from 100 μ L serum using a refractometer (model RHC-200ATC, Westover Scientific).

In addition to the automated blood analysis, thin blood smears were made from EDTA blood for manual differential cell counts. These smears were transported to the University of Pretoria, South Africa, where blood smears were stained with Diff Quick (Kyrion South Africa) for differential counts to calculate the absolute lymphocyte count (Lymph), absolute eosinophil count (Eos), absolute neutrophil count (Neut) and absolute monocyte count (Mono).

Descriptive statistics (mean and 95% confidence intervals) were used to describe the variation and spread in each hematological parameter of the calf population at each sampling point. Graphs are used to illustrate the changes in each parameter over time. The computation of the results and the production of the graphs were done with R 2.8.1 (Ihaka and Gentleman 1996).

Results

Figures 3-13 illustrate the changes for all parameters for the total calf population between day 7 and day 364.

The calves showed a significant increase in PCV, RCC and HGB from day 7 (means of 30%, $7.73 \times 10^6/\mu\text{L}$ and 10 g/dL, respectively) to day 42 (means of 36%, $10.23 \times 10^6/\mu\text{L}$ and 12 g/dL) (Figs. 3-5). From day 42, the general trend for all three parameters was a decrease until day 364. The RDW was relatively high between day 7 to 42 (means of 34.9 and 35.9 fL, respectively), indicating a significant variation in the size of red blood cells, which is suggestive of a high number of immature red blood cells (Fig. 6). From day 42 to day 112 the RDW decreased significantly, and then it remained between 31.2 and 31.8 fL to day 364. The MCV (Fig. 7) was relatively high at day 7 (mean 39.8 fL), confirming the presence of a high

number of immature red blood cells, but decreased gradually from day 7 up to day 147 (mean 33.6 fL), from where it increased gradually until day 364 (mean 36.0 fL). The MCHC (Fig. 8) showed a similar but more gradual trend, where it decreased between day 7 (mean 32.2 g/dL) and day 77 (mean 31.6 g/dL), and then increased up to day 364 (mean 33.05 g/dL).

The WCC and Lymph (Figs. 9 and 10) showed very gradual increases from day 7 (mean 9.00 $\times 10^3/\mu\text{L}$ and 3.7 $\times 10^3/\mu\text{L}$) to day 364 (mean 11.40 $\times 10^3/\mu\text{L}$ and 7.6 $\times 10^3/\mu\text{L}$). The Eos (Fig. 11) increased considerably from day 7 (mean 0.27 $\times 10^3/\mu\text{L}$) to day 322 (mean 0.75 $\times 10^3/\mu\text{L}$). The Neut showed a gradual decrease from day 7 to day 112, after which it remained between 2.4-2.55 $\times 10^3/\mu\text{L}$ (mean) until day 287 and increased again to day 364.

The Plt (Fig. 12) also showed a steady decrease from day 7 (mean 611 $\times 10^3/\mu\text{L}$) up to day 364 (mean 385.5 $\times 10^3/\mu\text{L}$). The MPV was relatively high at day 7 (mean 6.5 fL), decreased gradually up to day 112 (mean 5.8 fL) and increased up to day 364 (mean 6.2 fL).

The TSP (Fig. 13) was relatively high at day 7 (mean 9.8 g/dL), after which it decreased up to day 112 (mean 7.6 g/dL) but then remained between 7.6-8.0 g/dL (mean) up to day 364.

Discussion

This study showed that the hematological profile of the Short-horn Zebu differed both in levels and age-related trends compared to reference ranges reported for European breeds. For comparison, reported ranges for PCV for other indigenous cattle breeds are depicted in Table 1 and for European breed in Table 2, for which age-related changes are reported in more detail.

The most significant age-related changes in the red blood cell parameters of the Short-horn Zebu calves in this study occurred between day 7 and day 112. Concurrent low RCC and PCV with a relatively high MCV and RDW in the first week suggest a macrocytic anemia. A high

RDW value is an indication of anisocytosis which is often seen in regenerative responses due to a high number of reticulocytes. Following day 7 up to day 42 there was a significant increase in PCV, HGB and RCC accompanied by a high RDW, suggestive of increased erythropoiesis. The high MCV and a concurrent decrease in MCHC seen between day 7 and day 42 also indicates an increase in the number of immature RBC which is indicative of increased erythropoiesis. In neonatal Holstein calves erythrocytes were found to decrease in size up to the age of 3-4 months (Mohri et al. 2007), resulting in a gradual decrease in MCV. Immature red blood cells do not yet produce hemoglobin optimally since hemoglobin production per cell increases as red blood cells mature (Harvey 1989), hence a lower MCHC.

After day 42 there was a decrease in PCV and HGB together with a continuing decrease in MCV and MCHC, while RCC started decreasing from day 77. This may be suggestive of the development of microcytic hypochromic anemia. Iron supplementation during the first four weeks after birth corrected the drop in HGB in dairy calves (Mohri et al. 2004). The iron levels of calf serum were not evaluated in this study but could possibly have added value to explaining the results in this study.

The relatively low numbers of WCC and Lymph and high Neut around birth are consistent with what is reported for other cattle breeds (Knowles et al. 2000; Mohri et al. 2007) and are due to perinatal stress and high levels of cortisol during partus (Jain 1993). After the first week, levels in WCC, Lymph and Eos increased as the immune system matured and the animals became exposed to pathogens with resultant cellular immune responses. The Eos and Mono of the Short-horn Zebu were considerably higher than published ranges for neonatal calves of European breeds (Knowles et al. 2000; Mohri et al. 2007), but are comparable to ranges reported for other indigenous African cattle breeds (Oduye and Okunaiya 1971) and are possibly related to high parasite burdens under field conditions.

The platelet counts of the Short-horn Zebus were relatively high at day 7 and then decreased significantly up to day 364. The MPV was also relatively high at week 1. Together with a high RDW at day 7 this is suggestive of a regenerative response by the bone marrow at the time around birth in both red cells and platelets.

The initial high levels in TSP in the first week were likely due to the uptake of immunoglobulins from colostrum. The degradation of absorbed immunoglobulins likely contributed to the age-related decrease in TSP after the first week. Chronic parasitic infections, such as severe helminthosis, or malnutrition or a combination of both could possibly also have contributed to the gradual decrease in TSP after day 7.

The increase found in the red cell parameters during the neonatal period in the Short-horn Zebu calves in this study is in contrast to what is described for other ruminants, including European cattle breeds (Karesh et al. 1986; Knowles et al. 2000; Mohri et al. 2007). For these species and breeds there is a significant decrease in PCV, HGB and MCV in neonates up to around 6 weeks that coincides with an increase in RCC, after which all three parameters increase up to adult levels (Karesh et al. 1986; Knowles et al. 2000; Mohri et al. 2007) (Table 2). These changes, which are considered physiological, are ascribed to a decline in fetal erythrocytes at a faster rate than the production of adult-type erythrocytes (Karesh et al. 1986). This also coincides with the replacement of fetal hemoglobin by adult hemoglobin (Jain 1993; Mohri et al. 2007). Ruminants and primates have a distinct type of hemoglobin during fetal life (Jain 1993). In these species embryonal hemoglobin is soon replaced by fetal hemoglobin (Hb F), which in turn is eventually replaced by adult hemoglobin (Hb A). Hemoglobin F has a higher affinity for oxygen than Hb A and its function is to maintain partial pressure of oxygen of fetal blood. The replacement of Hb F with Hb A, referred as “switching”, occurs within the first few weeks after birth in ruminants.

In contrast to other ruminant and European cattle breeds, the replacement of fetal erythrocytes by adult-type cells evidently does not cause a decrease in HGB and PCV after birth in the Short-horn Zebu calves. Whether the rise in HGB and PCV seen in these calves coincides with the switching from Hb F to Hb A warrants further investigation. If Hb F replacement by Hb A is more rapid and efficient in reaching adequate Hb A levels before Hb F starts declining, or if there is a delay in the decline of Hb F while Hb A is produced by an increasing number of adult-type erythrocytes, such that the total HGB is maintained or even increased, it would be a possible explanation for the increase in HGB, PCV and RCC seen during the first weeks after birth for this breed.

Prolonged postnatal production of Hb F has been described in humans in certain hemoglobinopathies and thalasemias where production of Hb F continues into adulthood (Pasvol et al. 1976; Nagel 1990). It is considered benign, but has been shown to be protective against malaria in early childhood by retarding the intraerythrocytic growth of *Plasmodium falciparum* (Pasvol et al. 1976; Nagel 1990). In areas where malaria is endemic there is a high frequency in the genes for these hemoglobinopathies in the local human populations.

Prolonged production of Hb F in humans has also been described in neonates under hypoxic stress (Shiao and Ou 2006). The production of Hb F is also pharmacologically induced in the treatment of sickle-cell anemia.

Anaemia is an important hematological manifestation of most endemic blood-borne diseases of cattle in the tropics, including anaplasmosis, babesiosis and trypanosomosis. Western Kenya is an ECF endemic region and the calves in this study were shown to be exposed to and become infected with several of the tick-borne pathogens and trypanosomes from a very early age. East African Short-horn Zebus are considered to be disease resistant, however, particularly against tick-borne diseases. Further diagnostic tests are required to determine whether delayed switching between hemoglobin types can explain the postnatal rise in HGB in this breed, but it makes for interesting speculation whether prolonged postnatal production

of HbF plays any part in the innate resistance of Zebu calves against blood-borne pathogens during early calf-hood.

Compared to published ranges for European cattle breeds (Knowles et al. 2000; Mohri et al. 2007) TSP in these Short-horn Zebu calves was considerably elevated during the entire monitoring period, but is comparable to ranges reported for other African Zebu breeds (Useh et al. 2008). One possible explanation for higher TSP compared to European breeds is a high level of antigenic stimulation resulting in high globulin levels. Even from an early age, exposure to pathogens is considerable under field conditions in the tropical environment. Unfortunately only total serum proteins were investigated. Without distinguishing between different proteins, in particular albumin and the various immunoglobulins, it is difficult to come to any conclusions with regard to the levels and trends of TSP levels in the calves in this study.

Despite its economical importance as the main cattle breed not only in Kenya, but in all of eastern Africa, little is known about baseline values of health parameters of the East African Short-horn Zebu. There is a need to establish breed-specific reference ranges for blood parameters for this breed. It is evident from this study that baseline values differ with age, particularly in the neonate, and possibly contributes to the physiological adaptability to environmental constraints and disease resistance of this breed. When one compares the age-related changes in the red blood cell parameters for the calves in this study to the reported values for other cattle breeds (Knowles et al. 2000; Mohri et al. 2007), there are considerable differences in both the ranges for different age groups, as well as the trends in change over time. The reference ranges for European breeds were established in cattle in environments that are controlled for disease and nutrition. For this reason a direct comparison to the East African Short-horn Zebu in this study is problematic to interpret, but it is of value since the physiology of age-related changes in these breeds has been studied more extensively than in indigenous African breeds. The changes in the red blood cell parameters of the calves under

study, especially during the neonatal period are not explained by what is known about the physiology of other cattle breeds. This warrants further research into the dynamics of blood cell parameters of the East African Short-horn Zebu and probably other indigenous cattle breeds.

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Conflict of interest

There were no conflicts of interests of any kind in the study design, collection, analysis and interpretation of data, writing of the manuscript, or the decision to submit the manuscript for publication.

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List of figures

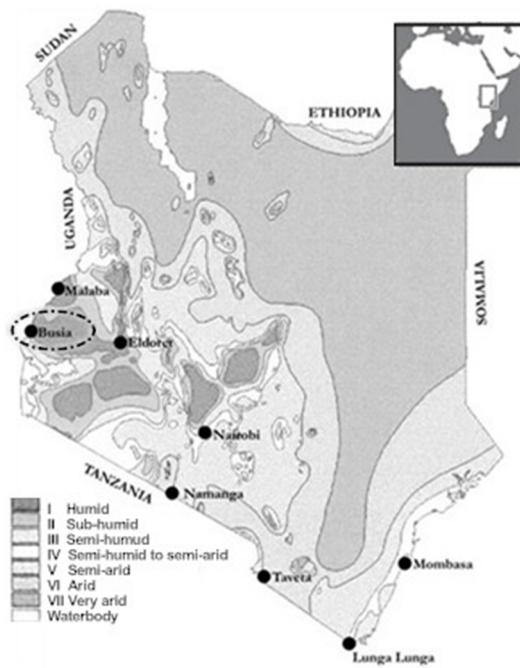


Fig. 1 Map of Kenya showing agroclimatic zones and highlighting the study area, Busia [adapted from map source: www.fao.org/docrep/008/y5968e/y5968e10.htm (2005)]



Fig. 2 An East African Short-horn Zebu calf at 51 weeks of age (69kg body mass)

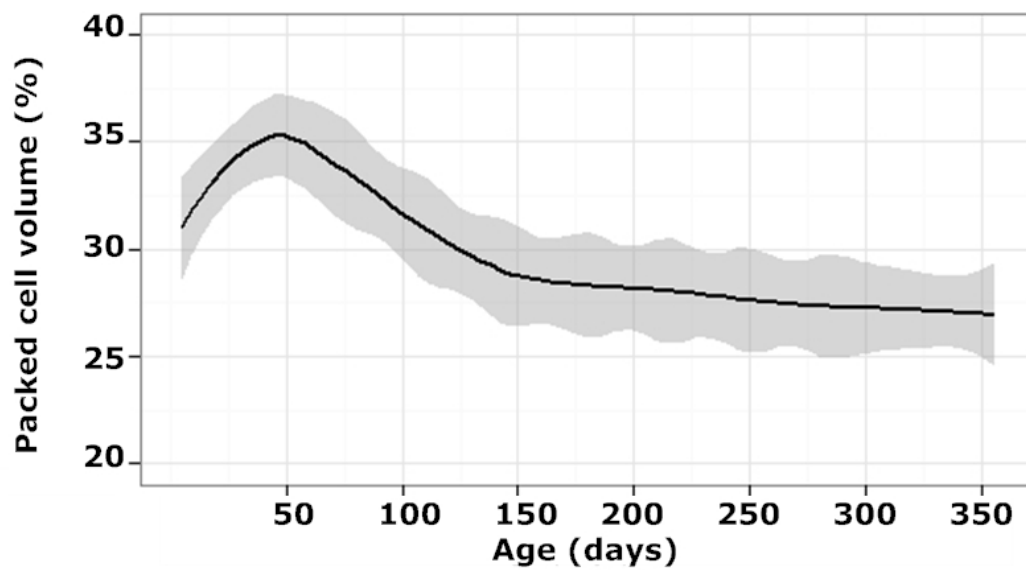


Fig. 3 Age-related changes in mean (95% CI) PCV of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

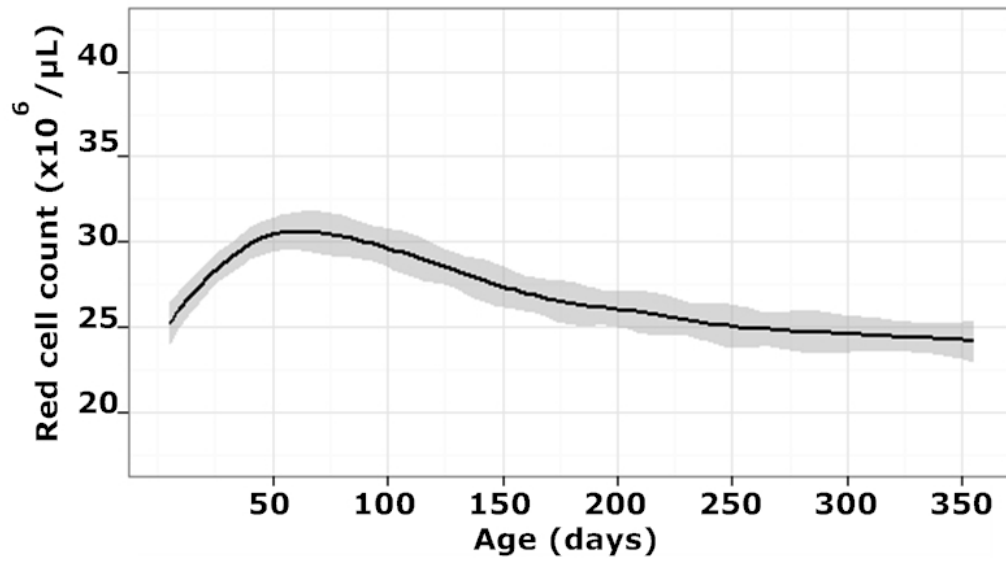


Fig. 4 Age-related changes in the mean (95% CI) RCC of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

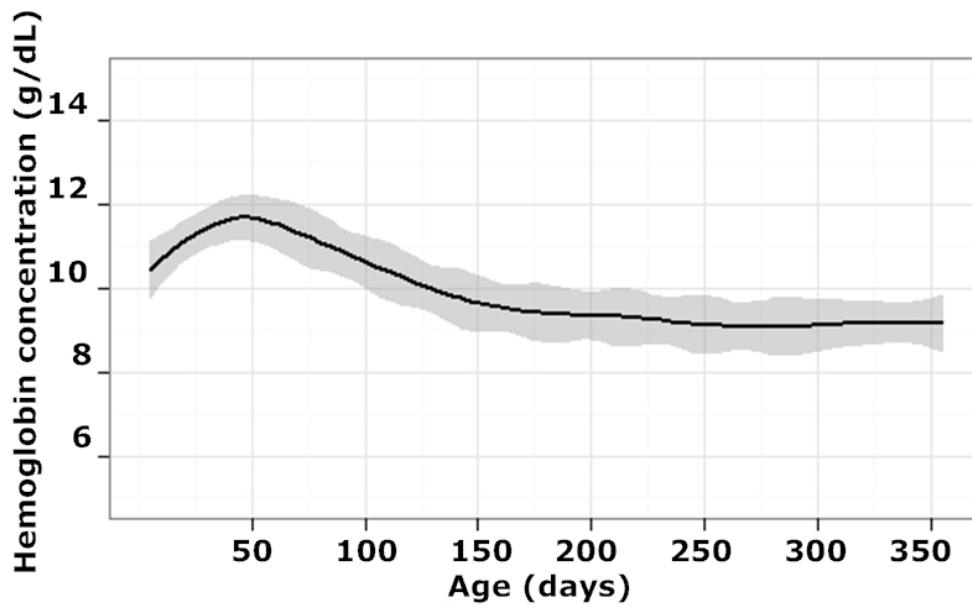


Fig. 5 Age-related changes in mean (95% CI) HGB of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

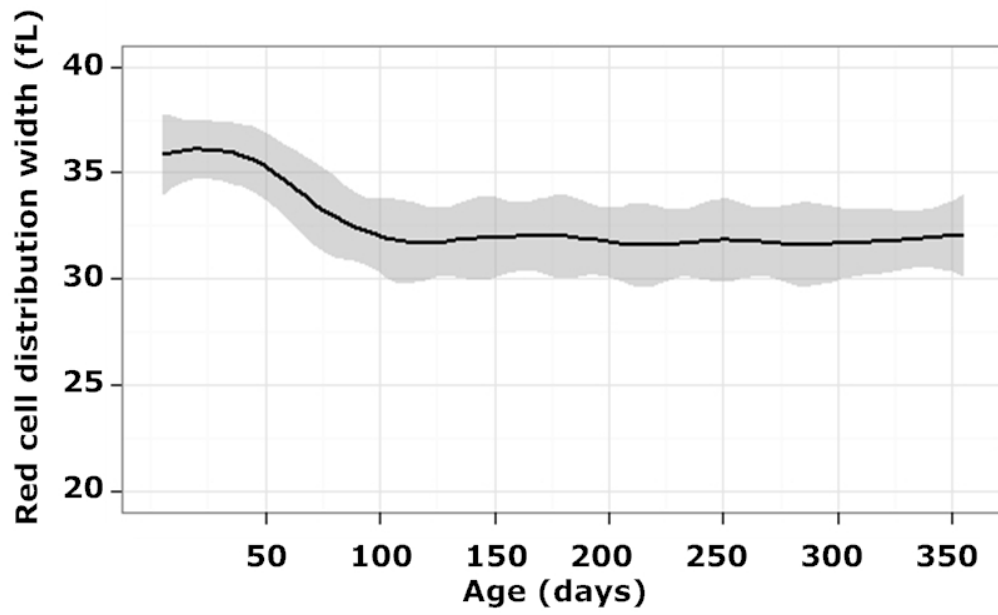


Fig. 6 Age-related changes in mean (95% CI) RDW of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

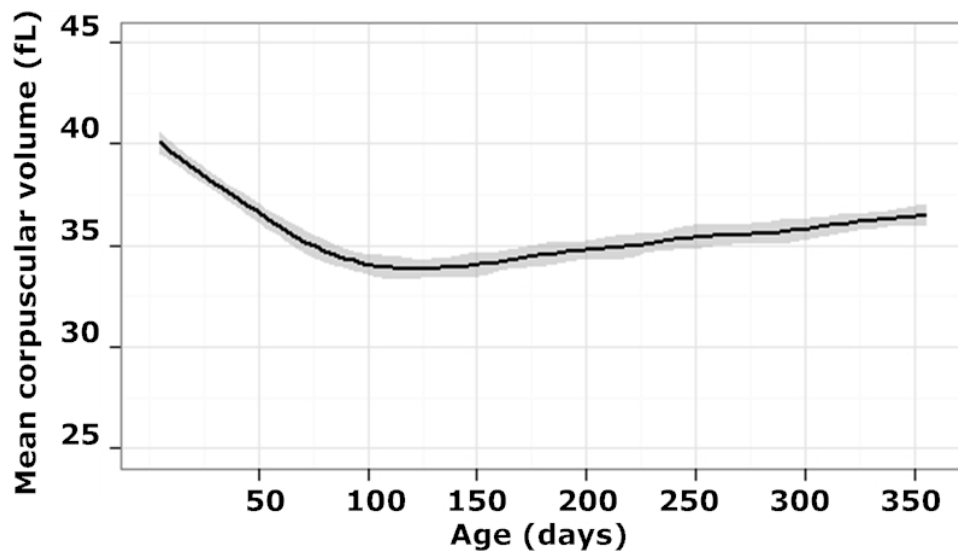


Fig. 7 Age-related changes in mean (95% CI) MCV of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

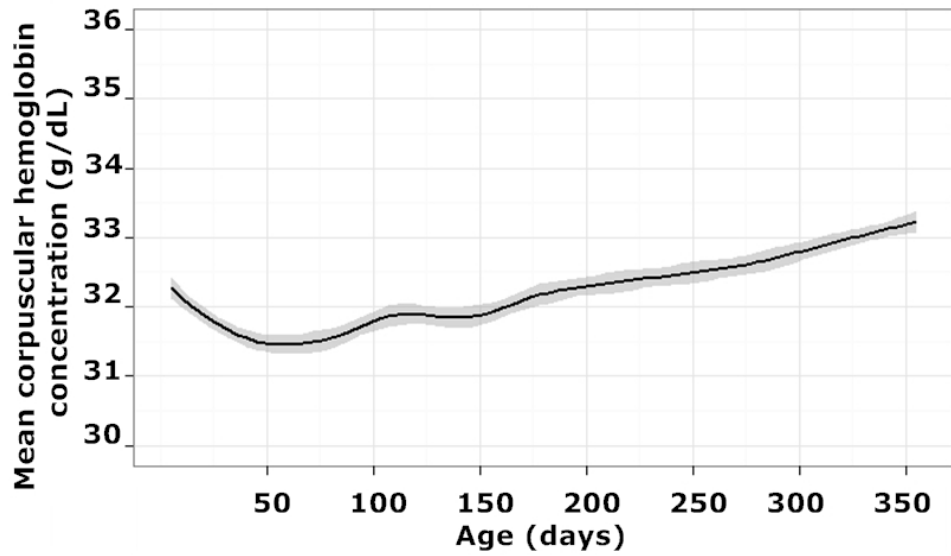


Fig. 8 Age-related changes in mean (95% CI) MCHC of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

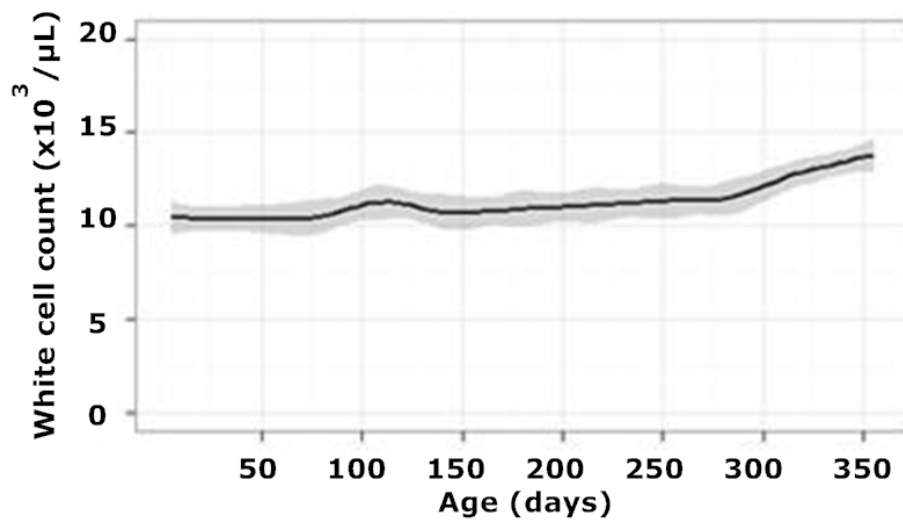


Fig. 9 Age-related changes in mean (95% CI) white cell counts of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

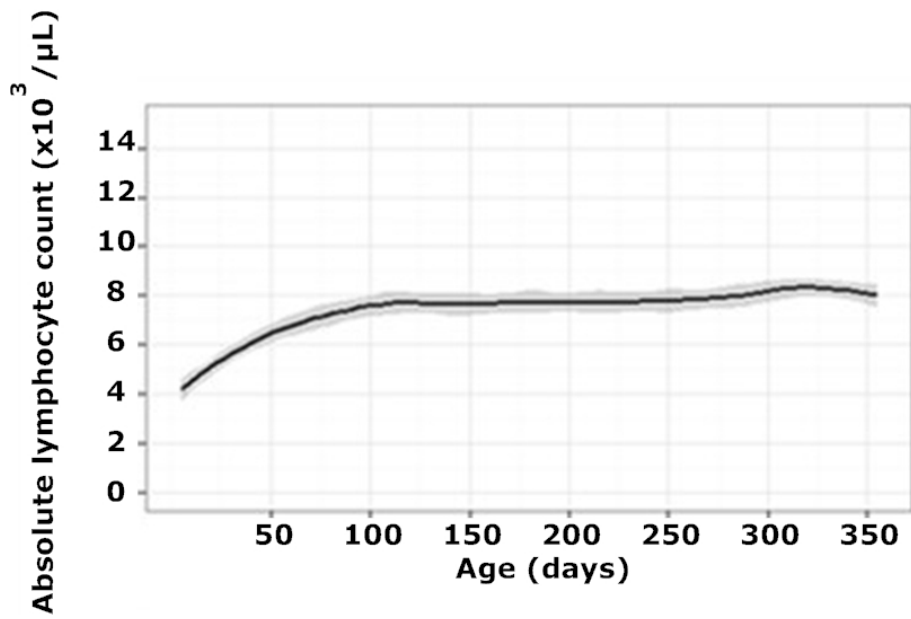


Fig. 10 Age-related changes in mean (95% CI) absolute lymphocyte counts of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

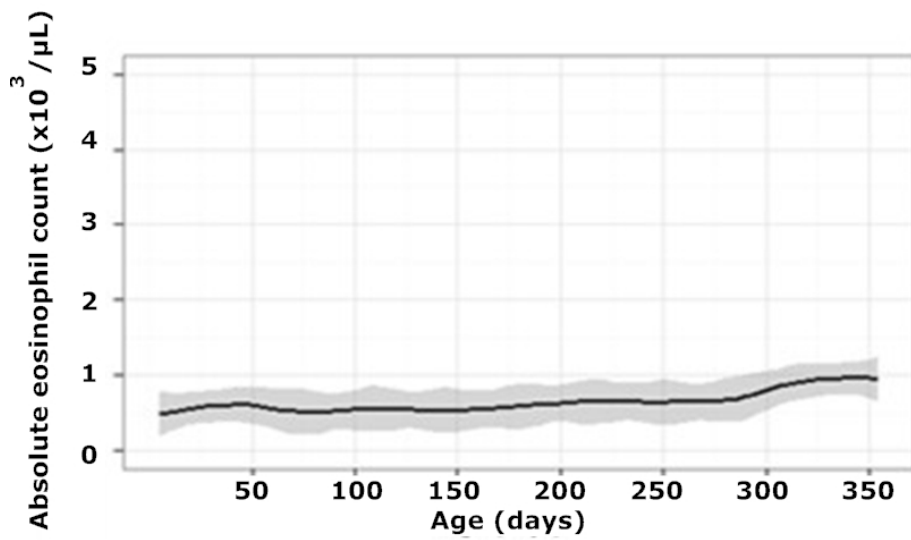


Fig. 11 Age-related changes in mean (95% CI) absolute eosinophil counts of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

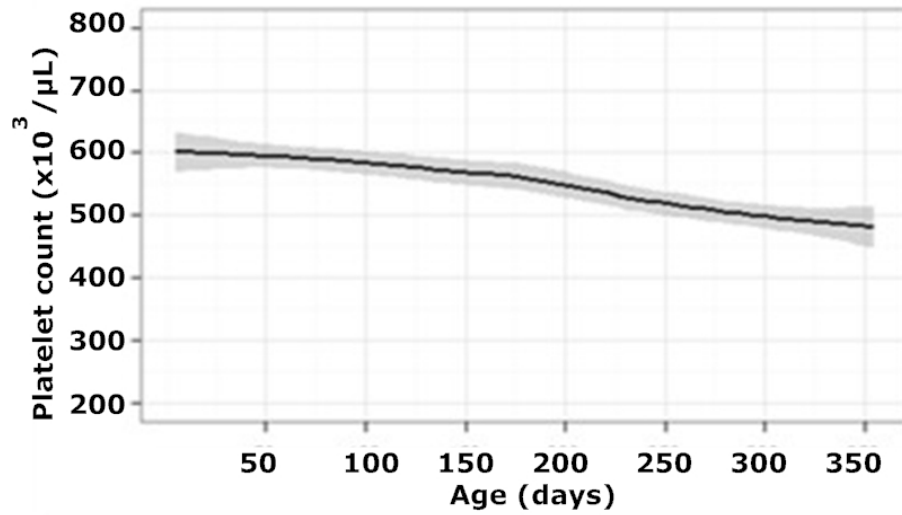


Fig. 12 Age-related changes in mean (95% CI) platelet counts of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

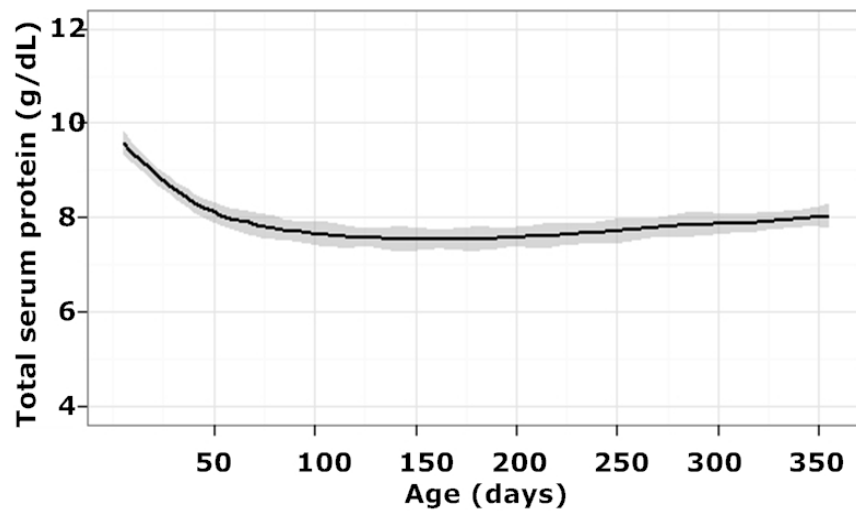


Fig. 13 Age-related changes in mean (95% CI) total serum protein of East African Short-horn Zebu calves from birth to 51-weeks of age (n=548)

List of tables

Table 1 Reported mean packed cell volume (%) of various African indigenous cattle breeds

Breed	Country	Age group	Mean PCV (%)	Reference
White Fulani (Zebu)	Nigeria	0-6 months	30.9 ¹	Oduye and Okunaiya (1971)
		7-12 months	28.5 ¹	
		13-24 months	29.1 ¹	
White Fulani	Nigeria	Adult	36.6 ± 5.5 (SD) ²	Saror and Coles (1973)
Zebu	Uganda	0-6 month	24.19 ± 1.9 (95%CI)	Magona et al. (2008)
		7-12 months	24.2 ± 1.5 (95%CI)	
		13-24 months	24.6 ± 1.1 (95%CI)	
Angoni	Zambia	Adult	29.1-31.8 ³	Van den Bossche and Rowlands (2001)
N'dama	Nigeria	N.S.	37.9 ± 3.6	Oduye and Okunaiya (1971)

¹ range not reported; ² during wet season; ³ range; N.S. not stated

Table 2 Reported mean packed cell volume (%) of various European cattle breeds

Breed	Country	Age group	Mean PCV (%)	Reference
Holstein	Canada	1 – 14 days	Range 17-47 ^a	Lumsden et al. (1980)
		2 weeks – 6 months	Range 23-42 ^a	
		6 months – 2 years	Range 26-48 ^a	
Holstein	Iran	1-2 days	27 ± 3 (SE)	adapted from Mohri et al. (2007)
		4 weeks	24 ± 3 (SE)	
		6 weeks	25 ± 3 (SE)	
		12 weeks	31 ± 2 (SE)	
Jersey	USA	3.5-4.5 months	36.16	Jain (1993)
		7.5-9 months	30.40	
		11-12 months	28.10	
Norwegian Red	Norway	1 week	35 ^a (range 25-39)	adapted from Brun-Hansen et al. (2006)
		6-8 weeks	30 ^a (range 19-37)	
		27-29 weeks	25 ^a (range 22-28)	

^a Hematocrit (L/dL)