

Evaluation of the hematological and biochemical changes of blood profile in Arabian horses under the effect of training and pregnancy

Fathia M. Benashour ^{1*}, Fahima A. Alnagar ², Nadia B. Gregni ³, Salheen M. Salheen ¹, Amal O. Buker ⁴ ¹ Department of Physiology, Biochemistry and Nutrition, Faculty of Veterinary Medicine, University of Tripoli, Tripoli, Libya. ² Department of Biochemistry and Molecular Biology, Faculty of Medicine, University of Tripoli, Tripoli, Libya. ³ Department of Statistics, Faculty of Science, University of Tripoli, Libya. ⁴ Department of Anatomy, Histology and Embryology, Faculty of Veterinary Medicine, University of Tripoli, Tripoli, Libya.

تقييم التغيرات الدموية والكيميائية الحيوية لصورة الدم في الخيول العربية تحت تأثير التدريب والحمل

⁴ فتحية محمود بن عاشور ¹*، فهيمة على النجار²، نادية البهلول القريقني³، صالحين محمد صالحين¹، أمال عمر بوكر⁴ ¹ قسم وظائف الاعضاء والكيمياء الحيوية والتغذية، كلية الطب البيطري، جامعة طرابلس، طرابلس، ليبيا. ² قسم الكيمياء الحيوية والاحياء الجزيئية، كلية الطب، جامعة طرابلس، طرابلس، ليبيا. ³ قسم الإحصاء، كلية العلوم، جامعة طرابلس، ليبيا. ⁴ قسم التشريح والأنسجة والأجنة، كلية الطب البيطري، جامعة طرابلس، طرابلس، ليبيا.

*Corresponding author: <u>fat.ashour@uot.edu.ly</u>

Received: April 29, 2024	Accepted: July 02, 2024	Published: July 13, 2024
h atua atu		

Abstract:

Knowing more information about Arabian horses raised in Libya is considered valuable before enrolling in any research on this category of horse's breed. This study investigated the influence of racing and pregnancy on the hematological and biochemical parameters. The blood samples were collected from 65 healthy Arabian horses in Libya for hematology and biochemistry analyses. The (WBC), (HB) and (HCT) were higher in racing and pregnant compared with (non-racing and non-pregnant) horses. The (MCV), (RDW-CV) was higher in pregnant than (non-racing and non-pregnant) horses. However, (MCHC) was significantly decreased in racing and pregnant animals. On the contrary, (MCH), (MPV), (PLT), (PDW) and (PCT) did not change among the examined Arabian horses. Biochemical analysis revealed a significant decrease in (ALT), (ALP), (AST) and (LDH), in racing and pregnant compared to the (non-racing and non-pregnant) horses with no significant different in AST. Although the (TP), and (IBIL) was significantly increased in race and pregnant compared to (non-racing and non-pregnant) horses, no changes were reported in (Alb), (T BIL), (D BIL) and (Glu) levels. The serum (TC) and (HDL) significantly decreased in racing and pregnant compared to non-pregnant animals with no significant change of the (TG), (VLDL), (LDL), (Cr) and urea of all animals. Finally, serum (P) and (K) were significantly lower, but (Mg) was significantly increased in racehorses and pregnant than non-pregnant mares and no changes were found in serum (Ca), (Na) and (Cl). It is crucial to understand these physiological changes to provide mares with the finest care and medical attention available. These data will help clinicians assess and care for pregnant mares appropriately.

Keywords: Arabian Horse, Hematology, Biochemistry, Racing, Pregnancy.

الملخص

إن معرفة المزيد من المعلومات عن الخيول العربية التي تتم تربيتها في ليبيا يعتبر أمرًا ذا قيمة قبل الالتحاق بأي بحث ضمن هذه الفئة من سلالات الخيول. بحثت هذه الدراسة في تأثير السباق والحمل على المعايير الدموية والكيميائية الحيوية. تم جمع عينات الدم من 65 حصاناً عربياً سليماً في ليبيا لتحليل معايير الدم والكيمياء الحيوية. كانت (WBC) و (HB) و (HCT) أعلى في خيول السباق والحوامل مقارنة بالخيول (غير المتسابقة وغير الحامل). وكان (MCV) و(RDW-CV) أعلى في الخيول الحامل منه عن (غير المتسابقة وغير الحامل) بينما انخفض مستوى (MCH) بشكل ملحوظ في حيوانات السباق والحوامل. وعلى العكس من ذلك لم تتغير (MCH)، (MCV)، (PDV)، (PDV)، و (PCP) بين الخيول (غير يشكل ملحوظ في حيوانات السباق والحوامل. وعلى العكس من ذلك لم تتغير (ALN) و (MCH)، (MCH)، والكام)، و (PCT) بين الخيول (غير العوامل مقارنة بالخيول (الحوامل. وعلى العكس من ذلك لم تتغير (ALN) و (ALN) و (ASN) و (LDN) في الخيول المتسابقة والحوامل العربية المفحوصة. أظهر التحليل الكيموحيوي وجود انخفاض مهم في (ALT) و (ALN) و (ASN) و (LDN) في الخيول المتسابقة والحوامل مقارنة بالخيول (غير المتسابقة وغير الحامل) مع عدم وجود اختلاف مهم في (AST). على الرغم من أن (PT)، و(IBII) زادا قياسيا في السباق والحوامل مقارنة بالخيول (غير المتسابقة وغير الحامل)، إلا أنه لم يتم الإبلاغ عن أي تغييرات في (ASN)، (CTBII) زادا قياسيا في السباق والحوامل مقارنة بالخيول (غير المتسابقة وغير الحامل)، إلا أنه لم يتم الإبلاغ عن أي تغييرات في (ASN)، (CTBII) زادا قياسيا في السباق انخفض قياسيا في العران ((GU)، (AST))، و(IDSI)، (IDSI) زادا قياسيا في السباق والحوامل مقارنة بالخيول (غير المتسابقة وغير الحامل)، إلا أنه لم يتم الإبلاغ عن أي تغييرات في (ASN)، (CTBII) زادا قياسيا في در (CT)، زادا تعايي في السباق انخفض قياسيا في مصل الذم (TC) و (TDSI) في السباقات والحوامل مقارنة بالحيوانات غير الحامل مع عدم وجود تغير معنوي في (TG)، (VLDI)، (VLDL)، (CT) واليوريا لجميع الحيوانات. وأخيرا، كان مصل الدم (P) و ((K))، وراح) ارتفع قياسيا في خيول السباق والحوامل مقارنة بالحيوانات غير (CSI)، زلال عالي والي في ديول السباق والحوامل مقارنة بالحيوانات غير الحامل مع عدم وجود تغير معنوي في خيول السباق (VLDI)، (VLDL)، (CT) واليوريا لجميع الحيوانات. وأخيرا، كان مصل الدم (P) و (CSI)، ورلكا)، ورلكا)، ورلكام العيوانات في مصل الدم (P) و ((CSI)، ورلكا)، ورلكا)، ورلكا، ورلكا)، ورلكا، ورلكا، ورلكا، ورلكا، مصل الدم (P) و (CSI)، ورلكا)، ورلكا، ورلكا

الكلمات المفتاحية: الحصان العربي، أمراض الدم، الكيمياء الحيوية، السباق، الحمل.

Introduction

Pregnancy has been demonstrated to cause hematological and biochemical abnormalities that may resemble illness patterns seen in people who are not pregnant. Hormonal milieu alterations are the primary cause of physiological changes during pregnancy [1, 2]. Pregnancy in many mammals, including mares, is associated with physiological changes that are reflected in hematological and biochemical profiles [2]. Although hormonal changes around the parturition have been well studied in the field of endocrinology [3], hematological and biochemical changes have been studied little [4].

Arabian horses are bred in stud farms where they live together with their conspecifics until they are 3 years old, and then they are taken to hippodromes where they are socially isolated from their conspecifics for flat races [5]. One of the most important roles of research in equine physiology is to obtain new useful information on characteristics that make the horse such a super athlete [6, 7]. In equine medicine, biochemistry and hematology are essential diagnostic techniques. It is crucial to distinguish these physiological changes from pathological ones that could endanger the health of the mare and/or fetus and necessitate rapid medical attention. These alterations have been connected to fetal development and are ascribed to the mares' health [8]. The hemogram and biochemistry are particularly important when working with sport horses subjected to heavy workloads. It allows tracking of the function of blood components, workability and performance of horses [9, 10]. It was reported that long-duration activities, such as endurance races, cause significant changes in hematological and biochemical components as a result of central and peripheral exhaustion [11]. A significant increase of 36% for the leukocytes (WBC, x10³/µL) was described in endurance horses immediately after exercise during the training season but no significant changes were noted for the erythrocytes (RBC, $x10^{6}/\mu$ L), hemoglobin (HB, g/dl) and hematocrit (HCT, %) [12]. After activity, deviations from the normal range in one or more blood components either above or below the range's boundaries might indicate a response to exhaustion by a working horse [11]. Moreover, an increase of RBC (19.1%), Hb (18.6%) and HCT (19.4%) were reported in race horses with increased Alanine aminotransferase (ALT. U/L) and total protein (TP, g/dl) significantly immediately after effort in endurance and race horses [12].

Previous studies have examined hematological and biochemical parameters in equine. these studies evaluated the blood values to provide normal reference intervals. To date, there is little information available about normal blood values during various phases of mare pregnancy or racing [13-15]. Several studies have reported hematology and biochemistry profiles in pregnant mares of different breeds including Spanish purebred [16], Carthusian [13], Holstein [17], and others [14, 18]. Thus, this study aimed to compare the hematological and biochemical values among three different horse physiological status (non-racing and non-pregnant, racing and pregnant mares).

Material and Methods

Animal welfare and ethical statements

All the experimental procedures and management were carried out in accordance with the Committee guidelines in the faculty of veterinary medicine, university of Tripoli, Tripoli, Libya. Regarding the protection of animals for experimental purpose and experimental conditions were accepted by institutional ethical committee.

Experimental animals

A total of sixty-five clinically healthy Arabian horses were used in this study at the winter season. The animals were grouped into (non-racing and non-pregnant) horses (n=34), racehorses (n=15) and pregnant mares (n=16) from one geographical area.

The animals were kept and fed by the owners and there were no changes in routine nutrition or husbandry. Prior to blood sampling, attending veterinarians ensured good health in all Arabian horses including assessment of medical history, physical examination, and routine anthelmintic and vaccine administration. If an animal showed any sign of disease, blood was not collected. Pregnancy dates were calculated based on the conception date.

Blood sampling

The blood samples were drawn from the left jugular vein using an 18-gauge needle and a 10-mL syringe then transferred into K₃EDTA. The samples from pregnant mares were collected during the last trimester of pregnancy (6 -10 months).

All specimens were properly marked with the subject identification number to avoid sample mix-ups. Blood serum was centrifuged at 3000 rpm for 20 min and subsequently stored at -20° C until the realization of biochemical analysis.

Hematological and biochemical analysis

All hematology and biochemistry analyses were performed by a reference laboratory (Esraa's and Alshefaa's clinical laboratories, Tripoli, Libya). The hematological indicators included in the study were as follow: leukocytes (WBC, $x10^3/\mu$ L), erythrocytes (RBC, $x10^6/\mu$ L), hemoglobin (Hb, g/dl), hematocrit (HCT, %), mean cell volume (MCV, fl), mean cell hemoglobin (MCH, pg), mean cell hemoglobin concentration (MCHC, %), red cell width distribution-coefficient of variation (RDW-CV, %), red cell distribution width-standard deviation (RDW-SD, fl), platelet distribution width (PDW, fl), and plateletcrit (PCT, %). The described indicators were examined with an automatic hematological analyzer (Celltac α , nihon kohden, tokyo, japan). platelets (PLT, $x10^3/\mu$ L), mean platelet volume (MPV, fl), platelet distribution

The serum biochemical indicators included in the study were as follow: alanine aminotransferase (ALT, U/L), alkaline phosphatase (ALP, U/L), aspartate aminotransferase (AST, U/L) and lactate dehydrogenase (LDH, U/L), glucose (Glu, mg/dl), total protein (TP, g/dl), albumin (Alb, g/dl), total and direct bilirubin (T BIL, D BIL, mg/dl),total cholesterol (TC, mg/dl), triglycerides (TG, mg/dl), very low density lipoprotein (VLDL, mg/dl), low density lipoprotein (LDL, mg/dl), high density lipoprotein (HDL, mg/dl), urea (mg/dl), creatinine (Cr, mg/dl), calcium (Ca, mg/dl), phosphorus (P, mg/dl), and magnesium (Mg, mg/dl) were measured using an automated blood chemistry analyzer (pz Cormay ACCENT M320) as well as serum sodium (Na, mg/dl), potassium (K, mg/dl), and chloride (Cl) were determined using easylyte plus automatic analyzer using ion-selective electrode. Indirect bilirubin (I BIL) was estimated as the arithmetical difference between serum total bilirubin and direct bilirubin values.

Statistical analysis

The data was submitted to statistical analysis using the computational program SAS version 25. Values were compared using ordinary one-way ANOVA method and the level of significance was set to: p < 0.5, p < 0.01, p < 0.001, p < 0.0001 and column statistics were calculated. For the non-normally distributed data, a Kruskal-Wallis analysis was used and when data were statistically significant.

Results and discussion

Table 1&2 summarizes the hematological indices of (non-racing and non-pregnant) horses, racehorses and pregnant mares. The WBC, Hb and RBC showed an increase in racing and pregnant mares but this did not significantly different in comparison with the (non-racing non-pregnant horses). However, the HCT values was significantly (p < 0.005) higher in racing and pregnant mares compared with (nonracing and non-pregnant) horses. Similar findings obtained by [19] who reported that WBC, HCT, HGB, PLT, RDW-SD, and RDW-CV were not significantly modulated (p > 0.05) in Thoroughbred racehorses. Also, [2, 20] stated that total WBC counts and differentials were not statistically significant between pregnant and non- pregnant mares. Physiologically, higher RBC, HCT and HGB levels can play a role in better physical condition, such as in sports activities [21]. Previous investigations displayed significantly lower values of RBC, HGB, and HCT in pregnant mares compared with non-pregnant mares [2, 20]. According to fetal growth that occurs in that period of pregnancy produces a greater oxygen demand [22]. This greater need for oxygen is compensated by the endocrine system that stimulates the release of erythropoietin by the renal tissue [23]. The secretion of this circulating glycoprotein stimulates increased production of erythrocytes in the bone marrow [24]. This phenomenon, which has been described in pregnant rabbits [25] and pregnant women [22], has also been observed in Arabian pregnant mares [20, 21]. Additionally, studies on rabbits stated an increase in RBC, HCT, and HGB during the 2nd trimester followed by a reduction in the 3rd trimester [26, 27]. A previous study on 16 draft mares reported no significant changes in RBC, HCT, and HGB before and

after delivery [4]. Moreover, there was no significant difference in the erythrocyte variables between pregnant and non-pregnant Campeiro mares [28] or pregnant and non-pregnant Warmblood mares [29] or between pregnant and non-pregnant Barb mares [30]

The MCV (p=0.01), RDW-CV (p<0.001) and RDW-CV (p<0.001) showed a significant higher value (p<0.01) in pregnant mares than non-racing non-pregnant horses. The same findings obtained by previous report [31] who found significantly higher values for the MCV in pregnant Crioula mares. The results found for MCV and RDW suggest an increase in the number of immature erythrocytes, which would be in accordance with findings in pregnant rats, rabbits and women reported by [20, 24, 25]. However, MCHC was significantly (p-<0.001) decreased in racing and pregnant animals. Decreased circulating lifespan of erythrocytes and hemodilution have been proposed as mechanisms for decreased RBC and HB concentrations during pregnancy in humans and rabbits. The MCH was non-statistically different among all animals. Our results partially agree with [2] who reported that MCV, MCH, and MCHC did not significantly change between pregnant and non-pregnant mares. Previous studies on pregnant mares reported no significant changes in MCV, and MCHC [4, 14, 32]. Similarly, no significant differences were established in MCV, MCH and MCHC in endurance horse's pre and post 120 km race [9]. However, [33] observed an increase in RBC count, Hb and HTC in endurance horses competing at a distance from 30 km to 100 km.

lterree	Dhur	Maan		Std.	Kruskal-Wallis test	
Items	Pnys.	wean		Deviation	test statistics	<i>p</i> -value
	no Race, no pregnant	6.82	±	1.92		
(v10 ³ /uL)	Race	6.77	±	1.31	0.657	0.720
(x10 /μΕ)	pregnant	7.07	±	1.24		
	no Race, no pregnant	12.06	±	2.13		
(a/dl)	Race	13.05	±	2.57	1.455	0.483
(3,)	pregnant	12.39	±	1.65		
	no Race, no pregnant	6.94	_ ±	1.03		
RBC (x10 ⁶ /µL)	Race	7.70	±	1.14	5.647	0.059
	pregnant	7.44	±	0.98		
HCT	no Race, no pregnant * (p-value < 0.005)	30.91	±	5.74	40.444	0.000
(%)	Race *	36.05	±	5.35	12.414	0.002
	Pregnant *	35.81	±	4.99		
MCV	no Race, no pregnant * (p-value =0.010)	44.48	±	3.82	0.005	0.010
(fl)	Race	46.99	±	3.43	0.225	0.016
	Pregnant *	48.46	±	4.61		
MOLL	no Race, no pregnant	17.07	±	1.23		
MCH (pg)	Race	16.38	±	1.79	1.543	0.462
(P9)	Pregnant	16.36	±	1.84		
МСНС	no Race, no pregnant * (p-value <0.004)	38.67	±	3.84		
(%)	Race*	34.88	±	2.65	16.912	0.000
	Pregnant*	33.91	±	1.27		
RDW-CV	no Race, no pregnant * (p-value <0.003)	18.41	±	2.89	-	
(fl)	Race *	21.52	±	2.71	15.891	0.000
	Pregnant *	21.60	±	2.61		
RDW-SD	no Race, no pregnant * (p-value <0.002)	28.42	±	9.26	40.500	0.000
(%)	Race*	37.81	±	6.76	16.523	0.000
	Pregnant*	38.51	±	6.88		

Table 1. Hematological values in non-racing non-pregnant horses (n=34), racehorses (n=15) and pregnant mares (n=16) (Mean ±SD).

* Indicate significant difference (p<0.05%) among the different animals.

Items	Dhue	Maan		Std.	Kruskal-Wallis test	
	Phys.	Mean		Deviation	test statistics	<i>p</i> -value
	no Race, no pregnant	81.96	±	49.08		
PLI (v103uL)	Race	82.01	±	50.44	1.328	0.515
(χτο με)	pregnant	73.72	±	60.20		
MPV	no Race, no pregnant * (p-value =0.02)	7.15	±	0.57	0.007	0.040*
(fl)	Race	7.40	±	0.43	6.297	0.043
	Pregnant *	7.57	±	0.57		
	no Race, no pregnant	15.16	±	1.38		
PDVV (fl)	Race	15.25	±	2.86	5.566	0.062
(1)	pregnant	15.97	±	2.14		
PCT (%)	no Race, no pregnant	0.07	±	0.09		
	Race	0.06	±	0.03	0.617	0.735
	pregnant	0.10	±	0.16		

 Table 2. Platelet count and platelet indices in non-racing non-pregnant horses (n=34), racehorses (n=15) and pregnant mares (n=16) (Mean ±SD).

* Indicate significant difference (p<0.05%) among the different animals.

Concerning platelets evaluations (Table 3), with the exception of mean platelet volume (MPV), PLT count, platelet distribution width (PDW) and plateletcrit (PCT) values reported no significant changes among the examined Arabian horses (Table.2). This agree with [4, 14, 32]. The MPV, PDW and PCT parameters are indicators of PLT, they indicate changes in the form and activation of platelets, and of them PDW is the most specific marker [34]. According to [35], the analysis of MPV changes may be a potential marker in pregnant cattle. As described in other studies [36], no statistically significant differences were found in the MCH, mean platelet volume (MPV), thrombocrit or red blood cell distribution width (RDW) in the pregnant Polish Holstein-Friesian black-and-white cattle. Also our results were parallel to those obtained by [37] who reported that pregnant women showed a statistically significant increase in mean platelet volume (MPV) and the other platelet parameters showed differences, but were not statistically significant.

With reference to the biochemical results in this research, a significant decrease in serum ALT, ALP, and LDH activities were recorded in racing (6.53±5.36, 104.83±32.50, 324.22±78.99 U/L) and pregnant animals (5.11±3.06, 140.34±50.48, 305.76±109.15 U/L) compared to the non- racing non-pregnant horses (8.48±3.90, 182.09±89.74, 382.72±96.25 U/L). Moreover, there is lower activity of AST but did not significantly different as shown in Table 3.

On contrary to our results, the greatest increase in serum enzyme activity post-race was for those indicators of muscular injury, that is, CK, ALT, and AST. These enzymes were all within a normal reference range pre-race and changed dramatically, beyond the normal reference range, during the race [38]. According to a study by [39] the magnitude of increase of both CK and AST is not related to fatigue or performance (e.g., speed). In horses, the most accepted blood indicators of muscle damage are LDH and AST activities [12]. Other authors stated that activity of hepatic parameters is a health state monitoring factor in all organisms. Increased activity of AST is usually a result of muscle or liver diseases [40]. The AST activity decreased in racing or pregnant horses thus may be connected with the highest muscle fatigue during these conditions. Besides, the CPK and AST values are related to muscle mass and composition [41]. Furthermore, plasma CK, AST, and LDH activities of the horses that were not assessed to be poor performers were within normal RIs [15].

Items	Phys.	Mean	lean	Std. Deviation	Kruskal -Wallis test	
					test statistics	<i>p</i> -value
ALT	no Race, no pregnant	8.48	±	3.90		0.007*
(U/L)	Race	6.53	±	5.36	9.893	
	pregnant	5.11	±	3.06		
AST	no Race, no pregnant	263.65	±	74.06	E 490	0.064
(U/L)	Race	254.09	±	101.04	5.469	0.064
	pregnant	231.71	±	102.47		
ALP	no Race, no pregnant	182.09	±	89.74	11.986	0.002*
(U/L)	Race	104.83	±	32.50		
	pregnant	140.34	±	50.48		
LDH (U/L)	no Race, no pregnant	382.72	±	96.25	10.000	0.000t
	Race	324.22	±	78.99	10.282	0.006*
	pregnant	305.75	±	109.15		

 Table 3. Serum enzymes in non-racing non-pregnant horses (n=34), racehorses (n=15) and pregnant mares (n=16) (Mean ±SD).

* Indicate significant difference (p<0.05%) among the different animals.

As illustrated in Table 4, serum total protein (TP) and indirect bilirubin (I BIL) was significantly (p<0.05) increased in race (7.19 g/dl, 0.42 mg/dl) and pregnant horses (7.41 g/dl, 0.30 mg/dl) in relation to non-racing non-pregnant horses (6.80 g/dl, 0.64 mg/dl) but there is no statistical changes in serum albumin (Alb), total and direct bilirubin (T BIL, D BIL) and glucose (Glu) levels in between the studied experimental animals.

The increase in TP in racing and pregnant mares was in agreement with [28] who recorded significantly higher (P<0.05) values of TP in pregnant mares than non-pregnant mares. According to [42], during pregnancy, there is usually a decrease in serum albumin concentrations and an increase in globulin concentrations, followed by a decrease during the postpartum period due to the production of colostrum. Increase in globulin concentration due to pregnancy may have contributed to the increase in total protein concentration in pregnant mares.

About serum Alb results, analogous study reported no gestational changes in serum albumin levels in purebred Spanish and Holstein mares [16-18]. On the other hand, other investigators specified that Alb significantly decreased during the 2nd trimester of pregnancy [2]. Also, plasma concentrations of total protein, and albumin were reduced (p<0.001) from mid-gestation [43]. A similar pattern has been reported in women during the 3rd trimester. It has been stated that the reduction in Alb levels causes a decrease in oncotic pressure, leading to hemodilution in pregnant women, thus predisposing them to physiological edematous states and pre-eclampsia [44]. Additionally, plasma Alb concentration decreases during pregnancy and hemodilution is the more likely explanation may be related to reduced synthesis or increased catabolism of Alb [45]. There are some disagreements in previous reports regarding bilirubin levels. Former results show lower levels of T BIL (p< 0.001), D BIL (p< 0.001), and I BIL (p< 0.001) in pregnant compared with non-pregnant mares [2]. However, [18] reported higher T BIL concentrations in mid and late gestation in mares while [32] found that T BIL and D BIL were significantly higher in prepartum mares than in non-pregnant mares. Lower levels of T BIL, I BIL, and D BIL have been observed in humans as well, likely due to hemodilution [46]. Presumably, alterations in albumin levels and hemodilution are linked to alterations in bilirubin levels. Similarly, research on pregnant monkeys revealed a drop in liver functioning indicators, such as decreased levels of total bilirubin [47]. The serum glucose concentration decreased significantly during the race, while the triglyceride concentration increased substantially [38], findings that are consistent with those of earlier studies [48, 49]. However, the glucose level of thoroughbred race horses showed significant increase (p < 0.05) [6] and this increases may be attributed to hyper activity of sympathetic system and adrenaline release which activate hepatic glycogenolysis [50].

Items	Phys.	iys. Mean		Std.	Kruskal -Wallis test	
				Deviation	test statistics	<i>p</i> -value
TP	no Race, no pregnant	6.80	±	0.98	0.750	0.000*
(g/dl)	Race	7.10	±	0.52	9.752	0.008
_	pregnant	7.41	±	0.38		
ALB	no Race, no pregnant	3.69	±	0.51	2.424	0.000
(g/dl)	Race	3.74	±	0.33	2.421	0.298
_	pregnant	3.60	±	0.30		
T BIL	no Race, no pregnant	0.55	±	0.41	E 40E	0.075
(mg/dl)	Race	0.93	±	0.67	5.165	0.075
_	pregnant	1.12	±	1.02		
D BIL	no Race, no pregnant	0.25	±	0.19		0.066
(mg/dl)	Race	0.39	±	0.27	5.424	
_	pregnant	0.42	±	0.28		
I Bill	no Race, no pregnant	0.64	±	0.64		
(mg/dl)	Race	0.42	±	0.31	8.307	0.016*
	pregnant	0.30	±	0.33		
Gluc (mg/dl)	no Race, no pregnant	64.90	±	14.22	0.050	0.700
	Race	65.21	±	16.31	0.652	0.722
	pregnant	65.99	±	11.33		

 Table 4. Serum biochemical profile in non-racing non-pregnant horses (n=34), racehorses (n=15) and pregnant mares (n=16) (Mean ±SD).

* Indicate significant difference (p<0.05%) among the different animals.

Serum lipid profile and renal biomarkers (Cr and urea) in the present study showed a significant (p<0.05) decrease in serum TC and HDL in racing (88.55 mg/dl, 46.01 mg/dl) and pregnant mares (92.71 mg/dl, 51.14 mg/dl) compared with non-pregnant animals (103.00 mg/dl, 55.82 mg/dl), while the other determined parameters included TG, VLDL, LDL, Cr and urea were insignificantly changed among all animals as demonstrated in Table 5. However, in our study TG increased in pregnant mares but not statistically different. Similar results were obtained by [51] who showed a significant increase in TC , and TG in late pregnant thoroughbred Arabian race mares. The authors attribute the increase in TG to high very low-density lipoprotein (VLDL) production in liver increase due to negative energy balance in pregnancy and hyperlipemia may occur. Additionally, pregnancy in the Spanish broodmare induced hyperlipidemia with hypertriglyceridemia and increased plasma concentrations of total, direct and indirect bilirubin (p<0.05) [52].

On contrary to our findings, [6] stated that the urea and creatinine of thoroughbred race horses showed significant increase (p<0.05). Other investigation stated that blood ALT, AST, ALP, albumin, TB, and TC, TG, Cr, Na, and inorganic phosphate increased significantly with endurance racing, whilst chloride, glucose, iron, and potassium decreased significantly compared to pre-race levels [38].

Items	Phys	Mean		Std Deviation	Kruskal -Wallis test	
nomo	T Hyo.	mouri		etal Deviation	test statistics	p-value
тс	no Race, no pregnant	103.00	±	22.40	0.750	0.024*
(mg/dl)	Race	88.55	±	15.82	6.750	0.034
	pregnant	92.71	±	15.71		
TG	no Race, no pregnant	35.15	±	13.15	1 624	
(mg/dl)	Race	30.53	±	14.69	1.024	0.444
	pregnant	40.75	±	29.22		
HDL D	no Race, no pregnant	55.82	±	11.32	44.007	0.004*
(mg/dl)	Race	46.01	±	6.05	11.207	0.004"
	pregnant	51.14	±	7.41		
VLDL	no Race, no pregnant	14.97	±	4.74	3.860	0.145
(mg/dl)	Race	13.13	±	3.18		
	pregnant	13.19	±	3.21		
LDL	no Race, no pregnant	33.10	±	10.81		0.126
(mg/dl)	Race	28.91	±	9.66	4.143	
	pregnant	27.24	±	9.97		
CREA	no Race, no pregnant	1.29	±	0.20		
(mg/dl)	Race	1.37	±	0.22	2.602	0.272
(3 -) _	pregnant	1.35	±	0.24		
UREA (mg/dl)	no Race, no pregnant	36.09	±	6.89	1 224	0.516
	Race	33.36	±	5.26	1.324	0.516
	pregnant	35.16	±	5.95		

Table 5. Serum lipid profile and renal biomarkers in non-racing non-pregnant horses (n=34),racehorses (n=15) and pregnant mares (n=16) (Mean ±SD).

* Indicate significant difference (p<0.05%) among the different animals.

Concerning the mineral analysis in the experimental animals, serum P (p= 0.001) and K (p= 0.001) were significantly lower, meanwhile serum Mg was significantly increased in racehorses and pregnant mares than in non-pregnant mares. In addition, serum Ca, Na and Cl showed non-significant changes as illustrated in Table 6. The lower phosphorus and potassium levels in pregnant mares compared to non-pregnant mares may be due to fetal mineral consumption [53].

Maintaining optimal levels of Ca, P, Na, K and Cl is of particular importance for the water-electrolyte balance of the cell, which favors the transmission of nerve impulses and muscle contractions [54]. Prolonged exertion during endurance races or pregnancy is invariably accompanied by loss of electrolytes through sweat, urine, and to some extent with faeces. The loss of electrolytes through sweat leads to fatigue and weakness in horses [9]. Similar results were reported by [30] who stated that compared to non-pregnant Barb mares, P levels significantly decreased during the first, second, and third trimesters of pregnancy (p<0.014), (p<0.002) and (p<0.0001), respectively. P and Ca are essential for several body functions, including bone and energy metabolism. Most fetal Ca and P are deposited in the last 2 months of pregnancy, implying that most skeletal growth occurs at that time [55].

Serum K was significantly decreased in racing and pregnant horses and also sodium and chloride decreased in racehorses comparing with non-racing non-pregnant animals but did not statistically different. As described by other studies, [56] and [57] found a significant decrease in sodium as well as potassium and chloride during an endurance race in Australia and USA, respectively. Horses are known to secrete hypertonic sweat and lose sodium, chloride, and potassium through perspiration [54]. Also, the low concentration of potassium in the present study is most likely due to a combination of sweat loss, renal loss, water intake related dilution, and expected alkalosis as reported by [38]. Other study revealed that plasma concentrations of Na, K, urea, Cr, TG and TC were significantly reduced (p<

0.001) during pregnancy [43]. Decreased K concentration after a ride is a common finding [57-59], associated with loss in the sweat and increased activity of the Na-K-ATPase pumps during recovery.

ltems	Phys	Mean	Std Deviation	Kruskal -Wallis test		
nomo	i nyo.	moan			<i>p</i> -value	
C A	no Race, no pregnant	11.76	1.31			
(mg/dl)	Race	12.01	0.50	0.792	0.673	
	pregnant	11.78	0.69			
Р _	no Race, no pregnant	4.82	1.23		0.001*	
(mg/dl)	Race	4.14	0.96	14.084		
	pregnant	3.44	1.13			
Mg (mg/dl)	no Race, no pregnant	1.78	0.30			
	Race	1.93	0.20	9.309	0.010*	
(3 / _	pregnant	2.03	0.23			
No –	no Race, no pregnant	133.04	4.93		0.273	
(mEq/L)	Race	134.08	3.44	2.595		
、 ・ / =	pregnant	135.26	4.31			
к –	no Race, no pregnant	4.86	0.53			
(mEq/L)	Race	4.56	0.65	13.965	0.001*	
	pregnant	4.20	0.57			
CI (mEq/L)	no Race, no pregnant	102.24	4.43			
	Race	103.69	3.44	1.563	0.458	
	pregnant	104.62	4.33			

Table 6. Serum mineral profile in (non-racing and) non-pregnant horses (n=34), racehorses (n=15)and pregnant mares (n=16) (Mean ±SD).

* Indicate significant difference (p<0.05%) among the different animals.

Conclusion

Understanding such physiological changes is imperative to providing the best care and medical treatment in mares. These data will assist clinicians in better evaluating and treating any problems in the animals either race horses or pregnant mares.

Acknowledgments

We wish to Acknowledge the contribution of members of (AI- mosawwama farm), Tripoli-Libya. We also wish to express our gratitude to Mr. Ayoub Almusrati, the owner and manager of the farm for his valuable cooperation with us.

References

1. Chandra, S., et al., *Physiological changes in hematological parameters during pregnancy.* Indian journal of hematology and blood transfusion, 2012. **28**: p. 144-146.

2. Faramarzi, B., L.J. Rich, and J. Wu, *Hematological and serum biochemical profile values in pregnant and non-pregnant mares.* Can J Vet Res, 2018. **82**(4): p. 287-293.

3. Berg, E., D. McNamara, and D. Keisler, *Endocrine profiles of periparturient mares and their foals.* Journal of animal science, 2007. **85**(7): p. 1660-1668.

4. Aoki, T. and M. Ishii, *Hematological and Biochemical Profiles in Peripartum Mares and Neonatal Foals (Heavy Draft Horse).* Journal of Equine Veterinary Science, 2012. **32**(3): p. 170-176.

5. DANIŞAN, S., et al., *The Effect of Different Management Systems and Racing on the Stress Level of Arabian Horses.* Livestock Studies, 2023. **63**(1): p. 11-20.

6. Hassan, H.Y., et al., *The Effect of Race in the Clinical, Hematological and Biochemical Biomarkers in Thoroughbred Horses.* Europe's Journal of Psychology, 2015. **11**(3).

7. Fazio, F., et al., *Training and haematochemical profile in Thoroughbreds and Standardbreds: A longitudinal study.* Livestock Science, 2011. **141**(2): p. 221-226.

8. Gagnon, A., et al., *RETIRED: Obstetrical Complications Associated With Abnormal Maternal Serum Markers Analytes.* Journal of Obstetrics and Gynaecology Canada, 2008. **30**(10): p. 918-932.

9. Vlaeva, R., S. Sabev, and Z. Ivanova, *Hematological parameters in endurance horses pre and post 120 km race.* Revista de Ciências Agroveterinárias, 2023. **22**(1): p. 72-77.

10. Adamu, L., et al., *Effects of race distance on physical, hematological and biochemical parameters of endurance horses.* American Journal of Animal and Veterinary Sciences, 2010. **5**(4): p. 244-248.

11. Wakil, Y., et al., *Physical assessment, Hematological and Serum Amyloid A levels Pre and Post exercise in Arabian Horses in Maiduguri and Jere, Borno State. Nigeria: Physical Assessment, Hematological and Serum Amyloid A Levels Pre and Post Exercise in Arabian Horses.* International Journal of Equine Science, 2022. **1**(1): p. 11–15-11–15.

12. Maśko, M., et al., *The physical activity-dependent hematological and biochemical changes in school horses in comparison to blood profiles in endurance and race horses.* Animals, 2021. **11**(4): p. 1128.

13. Satue, K., O. Blanco, and A. Munoz, *Age-related differences in the hematological profile of Andalusian broodmares of Carthusian strain.* Veterinarni Medicina, 2009. **54**(4): p. 175-182.

14. Harvey, J., et al., *Haematological findings in pregnant, postparturient and nursing mares.* Comparative Haematology International, 1994. **4**: p. 25-29.

15. Bos, A., E. Compagnie, and A. Lindner, *Effect of racing on blood variables in Standardbred horses*. Veterinary Clinical Pathology, 2018. **47**(4): p. 625-628.

16. Satué, K. and P. Montesinos, *Plasma biochemistry in pregnant Spanish purebred broodmares*. Comparative Clinical Pathology, 2013. **22**: p. 113-117.

17. Milinković-Tur, S., et al., *Concentrations of total proteins and albumins, and AST, ALT and GGT activities in the blood plasma of mares during pregnancy and early lactation.* Veterinarski arhiv, 2005. **75**(3): p. 195-202.

18. Harvey, J.W., et al., *Clinical biochemistry of pregnant and nursing mares.* Veterinary clinical pathology, 2005. **34**(3): p. 248-254.

19. Cappelli, K., et al., *Immune Response in Young Thoroughbred Racehorses under Training.* Animals, 2020. **10**(10): p. 1809.

20. Orozco, C.A.G., et al., *Hematological values and total protein of Brasileiro de Hipismo and Breton mares during pregnancy.* Ciência Rural, 2007. **37**: p. 1695-1700.

21. Vaz, B., et al., *Constituintes hematimétricos do sangue de éguas gestantes de raça árabe.* Veterinária Noticias, 2000. **6**(1): p. 51-55.

22. Souza, A.I., M. B Filho, and L.O. Ferreira, *Alterações hematológicas e gravidez.* Revista brasileira de hematologia e hemoterapia, 2002. **24**: p. 29-36.

23. Plaschka, S., et al., *Parámetros hematológicos y bioquímicos en el parto de la yegua de raza española.* Medicina Veterinaria, 1997. **14**(4): p. 205-209.

24. Lurie, S., *Changes in age distribution of erythrocytes during pregnancy: a longitudinal study.* Gynecologic and obstetric investigation, 1993. **36**(3): p. 141-144.

25. Kim, J.-C., et al., *Haematological changes during normal pregnancy in New Zealand White rabbits: a longitudinal study.* Comparative Clinical Pathology, 2002. **11**: p. 98-106.

26. Cetin, N., T. Bekyürek, and E. Cetin, *Effects of sex, pregnancy and season on some haematological and biochemical blood values in angora rabbits.* Scandinavian Journal of Laboratory animal science, 2009. **36**(2): p. 155-162.

27. Wells, M.Y., et al., *Changes in clinical pathology parameters during gestation in the New Zealand white rabbit.* Toxicologic pathology, 1999. **27**(3): p. 370-379.

28. Fonteque, J.H., et al., *Hematological profile, total plasma protein and fibrinogen concentrations of clinically healthy adult Campeiro horses.* Ciência Rural, 2015. **46**: p. 144-149.

29. Nagel, C., et al., *Changes in blood pressure, heart rate, and blood profile in mares during the last 3 months of gestation and the peripartum period.* Theriogenology, 2016. **86**(7): p. 1856-1864.

30. Chikhaoui, M., et al., *Comparing hematological and biochemical profiles of pregnant and non-pregnant barb mares raised in Tiaret, Algeria.* Iranian Journal of Veterinary Medicine, 2023. **17**(4): p. 309-320.

31. Veiga, A., et al., *Haematological values, plasma protein and fibrinogen values of Crioulo horse-their variation with sex, age and management.* 2006.

32. Mariella, J., et al., *Hematologic and biochemical profiles in Standardbred mares during peripartum.* Theriogenology, 2014. **81**(4): p. 526-534.

33. Teixeira-Neto, A.R., et al., *Do hematologic constituents really increase due to endurance exercise in horses?* Pesquisa veterinária brasileira, 2012. **32**: p. 951-956.

34. Jakobsen, C., et al., *Platelet function in preeclampsia–a systematic review and meta-analysis*. Platelets, 2019. **30**(5): p. 549-562.

35. Saeed, O., et al. *Impacts of heat stress on blood metabolic in different periods of lactation and pregnancy in Holstein cows.* in *IOP Conference Series: Earth and Environmental Science.* 2021. IOP Publishing.

36. Wrzecińska, M., et al., *Examination of the haematological profile of pregnant Polish Holstein-Friesian black-and-white cattle in the early stage.* J Vet Res, 2023. **67**(3): p. 415-425.

37. Maconi, M., S. Cardaropoli, and A.M. Cenci, *Platelet parameters in healthy and pathological pregnancy.* J Clin Lab Anal, 2012. **26**(1): p. 41-4.

38. Larsson, J., et al., *Physiological parameters of endurance horses pre-compared to post-race, correlated with performance: a two race study from Scandinavia.* International Scholarly Research Notices, 2013. **2013**.

39. Kerr, M. and D. Snow, *Plasma enzyme activities in endurance horses.* Equine exercise physiology, 1983: p. 432-440.

40. DeNotta, S.L. and T.J. Divers, *Clinical pathology in the adult sick horse: the gastrointestinal system and liver.* Veterinary Clinics: Equine Practice, 2020. **36**(1): p. 105-120.

41. Witkowska-Piłaszewicz, O., R. Pingwara, and A. Winnicka, *The effect of physical training on peripheral blood mononuclear cell ex vivo proliferation, differentiation, activity, and reactive oxygen species production in racehorses.* Antioxidants, 2020. **9**(11): p. 1155.

42. Lopes, S.d.A., et al., *Manual de patologia clínica veterinária.* Santa Maria: UFSM-Universidade Federal de Santa Maria, 2007.

43. Harewood, W.J., et al., *The effects of the menstrual cycle, pregnancy and early lactation on haematology and plasma biochemistry in the baboon (Papio hamadryas).* J Med Primatol, 2000. **29**(6): p. 415-20.

44. Zannat, M., A. Nessa, and S. Ferdousi, *Serum albumin in first and third trimester of pregnancy*. Dinajpur Med Col J, 2016. **9**(2): p. 216-220.

45. Frederiksen, M.C. *Physiologic changes in pregnancy and their effect on drug disposition*. in *Seminars in perinatology*. 2001. Elsevier.

46. Bacq, Y., et al., *Liver function tests in normal pregnancy: a prospective study of 103 pregnant women and 103 matched controls.* Hepatology, 1996. **23**(5): p. 1030-1034.

47. Ibáñez-Contreras, A., et al., *Changes in blood parameters in rhesus monkeys (Macaca mulatta) during the first trimester of gestation.* Journal of Medical Primatology, 2013. **42**(4): p. 171-176.

48. Snow, D., et al., Alterations in blood, sweat, urine and muscle composition during prolonged exercise in the horse. 1982.

49. Spangfors, P., *Energibalans. vätske-och elektrolytförändringar under distansritt.* Svensk Veterinär Tidning, 1991. **42**(2): p. 55-61.

50. Trilk, J., et al., *A lactate-guided conditioning programme to improve endurance performance.* Equine Veterinary Journal, 2002. **34**(S34): p. 122-125.

51. Altiner, A., Changes in selected blood serum indices before and after parturition in mares. 2021.

52. Satué, K., et al., *Successful Pregnancy Outcome in Mares: The Potential Role of Body Conditional Score, Age and Biochemical Parameter's Adjustments.* Journal of Equine Veterinary Science, 2022. **115**: p. 104023.

53. Bazzano, M., et al., *Metabolic profile of broodmares during late pregnancy and early post-partum.* Reproduction in Domestic Animals, 2014. **49**(6): p. 947-953.

54. Thrall, M.A., et al., Veterinary hematology and clinical chemistry. 2012: John Wiley & Sons.

55. Kavazis, A., J. Kivipelto, and E. Ott, *Supplementation of broodmares with copper, zinc, iron, manganese, cobalt, iodine, and selenium.* Journal of equine veterinary science, 2002. **22**(10): p. 460-464.

56. ROSE, R.J., et al., *Plasma and sweat electrolyte concentrations in the horse during long distance exercise.* Equine Veterinary Journal, 1980. **12**(1): p. 19-22.

57. Schott, H., et al., *Body weight, fluid, electrolyte, and hormonal changes in horses competing in 50-and 100-mile endurance rides.* American Journal of Veterinary Research, 1997. **58**: p. 303-309.

58. Barton, M.H., et al., *Body weight, hematologic findings, and serum and plasma biochemical findings of horses competing in a 48-, 83-, or 159-km endurance ride under similar terrain and weather conditions.* American journal of veterinary research, 2003. **64**(6): p. 746-753.

59. Lindinger, M. and G.L. ECKER, *Ion and water losses from body fluids during a 163 km endurance ride.* Equine Veterinary Journal, 1995. **27**(S18): p. 314-322.