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# Field trials for Treatment of She Camels Mastitis in Sharkia Governorate

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### Abstract

Thirty mastitic milk samples were collected from she camels for the isolation and identification of bacteria causing mastitis and to determine their antibiogram against certain antibiotics. Bacteriological examination of mastitic milk samples revealed 18 single isolates (60%) and 12 mixed isolates (40%). Gentamicin was found to be the highest effective drug against the isolated bacteria than other used drugs. A total of 25 she camels (15 healthy and 10 mastitic) were divided into 5 equal groups receiving gentamicin alone and/or in combination with isoflupredone acetate. Blood and milk samples were taken from all she camels at 1<sup>st</sup>, 7<sup>th</sup> and 15<sup>th</sup> days post treatment for haematological and biochemical analysis. The results revealed a significant decrease in RBCs count, Hb, PCV%, serum total protein, albumin, globulin, Ca, Ph, Na levels in healthy she camels received gentamicin. While, isoflupredone acetate induced significant increase of WBCs count, AST, ALT, ALP, Ca, Ph, K levels in healthy she camels. Mastitis in she camels lead to significant decrease in RBCs count, Hb, PCV%, albumin, A/G ratio, Ca, Ph, Na, zinc, iron levels and milk production beside no statically difference in K and copper, WBCs, total protein, globulin, AST, ALT and ALP. Hematological and biochemical parameters alterations were returned to nearly normal levels on 10<sup>th</sup> day post treatment. It could be concluded that gentamicin and isoflupredone acetate had better results in reducing clinical signs of mastitis and improve adverse effects in she camels.

Keywords: Mastitis, She Camels, Gentamicin, Isoflupredone

# Introduction

Camels are most capable animal in utilizing marginal areas and in survival and production under harsh environment [1]. Camels are a good source of meat, milk, wool and hair [2]. Milk is synthesized in mammary gland [3]. Mastitis affects all domestic animals [4]. It has different causes and intensity degrees [5]. Udder infections originate from lymphatogenous or cutaneous routes [6]. It is caused by single or mixed infection [7]. It is infrequent in animals due to hand milking [8]. Gentamicin is an aminoglycoside antibiotic acting by inhibiting bacterial protein synthesis [9]. It has good efficacy in treating drug resistant G-ve bacteria [10]. Antiinflammatory drugs are widely used in veterinary practice to provide symptomatic relief of acute and chronic inflammatory conditions [11].

Isoflupredone acetate is a synthetic corticosteroid anti-inflammatory used in Veterinary Medicine. It is used as supportive treatment with antibiotics [12]. Combination of isoflupredone acetate with antibiotics has been approved by FDA [13]. No antagonistic interactions occurred between antimicrobial and anti-inflammatory activities [14].

The present study aimed to isolate and identify bacterial agents causing mastitis in she camel together with an antibiogram for the isolated strains, to evaluate the effect of mastitis on hemato-biochemical parameters in she camels.

#### **Material and Methods**

#### Drugs

Gentamicin (Garamycin<sup>®</sup>), Memphis Co. intramuscular injection. Each ml of solution

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contains 40-mg Gentamicin sulfate. Isoflupredone acetate (Predef  $2x^{\text{(B)}}$ ) a sterile injectable solution from Upjohn Co, Kalamazoo, U.S.A available as 50 ml vials.

### Animals and Experimental design

A total of 25 dairy she camels of 5-7 years old (15 healthy and 10 mastitic) from different localities in Sharkia Province were divided into 5 equal groups (Gp). Gp. (1) healthy she camels (control) Gp. (2) healthy she camels received 5 mg gentamicin /kg bwt. daily by I/M route for 4 days, Gp (3) healthy she camels received 0.2mg isoflupredone acetate/kg/B.wt daily by I/M route for 4 days, Gp. (4) mastitic she camels treated by gentamicin in the same doses and period, Gp. (5) mastitic she camels treated with gentamicin plus isoflupredone acetate in same dose, route and period.

### Milk samples and Bacteriological examination

Milk samples were taken from affected quarter in sterile bottles for bacteriological and chemical examination. Udder was washed with running water and dried with clean towel. Teats orifices were disinfected by 70% ethyl alcohol, after that few squirts of milk were discarded then milk sample from each infected quarter were collected. Milk samples were activated by incubation for 12h at 37oC then centrifuged at 3000 rpm for 30min. A loopful of sediment from each sample was streaked on surface of nutrient agar, MacConkey agar and blood agar containing 7.5% defibrinated sheep blood. plates were observed after All incubation for 24 h. at 37oC and any growth was recorded. Biochemical tests of isolated strains were performed [15].

# Antibacterial sensitivity tests

All isolated bacteria were used to check their susceptibility for gentamicin (10ug), cefotaxime (75ug), enrofloxacin (10ug), flumoquine (30ug), chloramphenicol (30ug), and oxytetracycline (30ug) [16].

# **Blood** samples

Two blood samples were collected from all she camels at 1<sup>st</sup>, 7<sup>th</sup> and 15<sup>th</sup> day post treatment, 1<sup>st</sup> sample was taken in a tube containing EDTA for studying blood picture [17] and 2<sup>nd</sup> sample was taken to obtain clear serum for estimation of total protein [18] albumin [19], transaminases (AST-ALT) [20] and ALP [21]. Serum and milk **c**alcium were determined [22], sodium and potassium were measured using flame photometer [23] inorganic phosphorus [24], copper[25], iron [26], zinc [27].

# Milk yield per day

Daily milk yield from healthy and mastitic she camels were collected before and at 1, 7 and 15 days post treatment.

# Statistical analysis

The obtained data were analyzed by T test [28].

# **Results and Discussion**

Mastitis is a serious inflammatory disease; its main clinical signs are depression, anorexia, fever, hotness beside pain of affected udder, swelling, firmness of infected quarters and clots in milk and in some cases milk become viscous. The same clinical signs were previously observed in mastitic she camels [29].

The etiological agents for mastitis in our study were single isolates (18) 60% (S. aureus (6) 20%, Corynebacterium spp. (3) 10%, E. coli (6) 20% and Streptococcus spp. (3)10%), mixed isolates (12) 40% (Corynebacterium +Streptococcus (3) spp spp 10%. Streptococcus spp + S. aureus (3) 10% S. aureus + E. coli (6) 20%) (Table 1). Identified bacteria isolated from mastitic milk samples were not re-isolated at 7<sup>th</sup> day post treatment. This result was similar to these reported by Obied et al. [30] in mastitic she camel. Tibary et al. [31] isolate Streptococcus spp. and E. coli from she camel mastitic milk and Bakeer et al. [32] isolate S. aureus from mastitic she camel.

No. of she camels	Type of isolates	No	%	Isolates	No	%	Re-isolation at 7 <sup>th</sup> day post treatment		
				S. aureus	6	20	– ve		
	Single			Corynebacterium spp.	3	10	– ve		
	isolate	18	60	E. coli	6	20	– ve		
30				Streptococcus spp.	3	10	– ve		
				Streptococcus +	3	10	– ve		
	Mixed	12	40	Corynebacterium					
	isolate			Streptococcus + S. aureus	3	10	– ve		
				$\tilde{S}$ . aureus + E. coli	6	20	– ve		

Table 1: Main bacterial isolates from mastitic milk causing mastitis in she camels (N=10)

The most effective antibiotic on isolated bacteria was gentamicin (Table 2). This result was similar with Abdel-Khalek and El -Sherbini They conducted [33]. that gentamicin had high effect on E. coli, S. Corynebacterium aureus, spp. and Streptococcus spp. Moreover Al-Juboori et al. [34] reported that, Streptococcus spp. isolated from mastitic milk of she camels was sensitive to gentamicin.

Mastitic she camels treated with gentamicin alone or plus Isoflupredone acetate revealed cure rate was of 100 % at 5 and 3 days post treatment respectively (Table 3). These results agreed with EL- Sheikh [35]. Treatment of mastitis in she camels with gentamicin plus isoflupredone acetate is one of most important potent treatment due to anti-inflammatory, anti-shock, anti-allergic and antitoxic activities of isoflupredone acetate as corticosteroid drug [36].

The obtained results revealed significant decrease in RBCs count, Hb, PCV% and

significant increase in WBCs count in healthy she camels receiving gentamicin, isoflupredone acetate or mastitic one (Table 4). Same results were previously recorded by Priuska and Schacht [37] they recorded that gentamicin chelates with iron and interfere with hemoglobin biosynthesis and formation of RBCs. These results are comparable with the results obtained by Naeshiro et al. [38]. They stated that gentamicin induces anemia due to deficiency of erythropoietin post kidney injury by gentamicin, site of erythropoietin production. Same haematological changes were previously recorded [39] in healthy sheep receiving Anti-inflammatory Isoflupredone acetate. drugs induce deleterious effect on bone marrow and changes in hemogram [40]. The same results were recorded in mastitic she camel [41]. Change in blood picture in mastitic animals may be due to inflammatory reactions [42], damage of erythrocyte by bacterial toxin [43].

Table 2: Sensitivity tests of isolated organisms against different antimicrobial agent

Antibiotic disc	Disc		Bacterial isolates								
Antibiotic disc	conc.	S. aureus	E. coli	Corynebacterium	Streptococcus						
Gentamicin	10ug	++++	+++	+++	+++						
Cefotaxime	75ug	+++	+++	++	++						
Enrofloxacin	10ug	++	+	++	++						
Flumequine	30ug	++	++	++	++						
Chloramphenicol	30ug	+	-	+	-						
Oxyteracycline	30ug	+	-	-	+						

Data presented in Table (4) revealed that gentamicin induce significant decrease in total protein, albumin and globulin but isoflupredone induce significant acetate decrease in T. protein, globulin and nonsnificantig decrease in albumin in healthy she camels, meanwhile mastitic she camel showed significant increase in total protein, globulin beside significant decrease in albumin and A/G ratio. Same changes were recorded in healthy sheep received gentamicin [44]. Reduction in protein picture may be due to damage of liver cells induced by gentamicin

[45]. The same reduction in protein picture post isoflupredone acetate treatment in rabbits was recorded by Nabila [46]. These changes protein picture may be due in to immunosupressive effect glucocorticoids [47]. Elevation in protein picture in mastitic she camels was recorded [29]. Reduction in serum albumin may be due to infiltration from blood to milk due to increase permeability of blood vessels as a result of inflammation [48]and/or due to damage of hepatic tissues by bacterial toxins [42].

Drugs	No. of she	3 days post treatment		5 days post treat	ment	6 days post treatment				
	camel	No. of cured animal	%	No. of cured animal	%	No. of cured animal	%			
Gentamicin	5	3	60	4	80	5	100			
G+I	5	5	100	-	-	-	-			

G+I: Gentamicin plus Isoflupredone acetate

The obtained results showed a significant increase in AST, ALT and ALP activities in healthy and mastitic she camels receiving gentamicin or isoflupredone acetate (Table 4). These results are in accordance with Sandhya and Varalakshmi [49] in healthy rats receiving gentamicin. Elevated liver enzymes induced by gentamicin may be due to liver damage [50]. Liver enzymes were elevated after isoflupredone acetate use in small animal [51]. Mastitis induced significant elevation in AST, ALT and ALP [52] and these elevations may be due to damage of hepatic tissues by bacterial toxins [53]

In the current work obtained data indicated nonsnificantig decrease in milk macro and micro elements, nonsnificantig increase in milk production post gentamicin or isoflupredone acetate administration to healthy she camel, but mastitis ones showed nonsnificantig decrease in milk Ca, Ph, K, micro elements and nonsnificantig increase in significant decrease in milk Na beside production (Table 5). Aminoglycoside antibiotic induce nonsnificantig decrease in milk copper, iron and zinc [54]. Isoflupredone acetate had no effect in milk production in healthy cows [55]. The same change in milk mineral was recorded by Bruckmaier *et al.* [56] in mastitic she camels and Batavani *et al.* [57] in mastitic cattle milk. Reduction in mastitic animals depended on degree of inflammation [58]. Mastitic cows treated with isoflupredone acetate and antibiotic showed an increase in production of milk [59].

Serum Ca, Ph, Na but K and microelements in healthy she camels received gentamicin were nonsnificantigly decreased but healthy she camels received isoflupredone acetate showed significant increase in Ca, Ph, nonsnificantig decrease in Na, microelements and K significantly decreased. Meanwhile, she camels showed significant Mastitic decrease in Ca, Ph, Na, zinc, iron, nonsnificantig decrease of K and cupper 5).Serum (Table macro elements were decreased healthy sheep received in gentamicin [44]. Similar results were also reported in rabbits injected with [60]. dexamethasone Mastitis induces reduction in serum mineral due to anorexic condition and decreased intestinal absorption of mineral [61]. Mastitis induced reduction in serum mineral may be due to anorexia in mastitic animals [62].

The hemato-biochemical parameters and milk elements in mastitic she camels were improved towards the normal level at  $15^{\text{th}}$  day post treatment with either gentamicin or plus Isoflupredone acetate. These results are in agreement with Hussein *et al.* [63] who reported that erythrogram and biochemical parameters in mastitic buffaloes were improved at  $10^{\text{th}}$  day post treatment with gentamicin.

#### Conclusion

It could be concluded that, mastitis induce several adverse effects on haemogram, biochemical parameters and milk elements which returned to the normal levels 15<sup>th</sup> days post treatment with gentamicin plus Isoflupredone acetate.

#### **Conflict of Interest**

The authors declare no conflict of interest.

		Dow	matan		Blood	picture			Protein pro	ofile (g/dl)		Liv	er enzymes (I	U.L)
Groups	Parameter			(X106/μl)	Hb (g/dl)	PCV %	WBC (X103/µl)	T.Protein	Albumin	Globulin	A/G	AST	ALT	ALP
Healthy	No	n treated (cont	rol)	9.83±0.90	11.37±0.90	29.39±0.93	13.41±0.73	8.45±0.30	4.74±0.21	3.71±0.50	1.28±0.21	44.13±1.94	23.07±0.88	36.26±1.96
she camels		Gentamicin		7.58±0.26*	8.13±0.16*	26.13±0.81*	14.46±0.84*	6.38±0.76*	3.30±0.3*	2.58±0.20*	$1.47 \pm 0.32$	49.20±1.12*	26.59±1.20*	43.05±1.25*
	Isof	flupredone ace	tate	7.37±0.34*	8.41±0.43*	26.42±0.69*	15.67±0.67*	6.13±0.46*	3.83±0.33	2.30±0.31*	$1.66 \pm 0.22$	48.96±0.37*	26.58±1.16*	43.86±1.40*
		pre treatment		7.23±0.31*	7.96±0.97*	26.97±0.99*	16.32±0.75*	9.17±0.13*	3.76±0.35*	5.49±0.36*	$0.68 \pm 0.12*$	50.26±1.67*	27.06±1.51*	43.65±1.47*
		Gentamicin	1st	7.52±0.15*	8.28±0.84*	26.06±0.96*	15.13±0.54*	9.17±0.11*	3.90±0.2*	5.27±0.30*	$0.74 \pm 0.10^{*}$	49.33±1.01*	28.16±1.60*	42.31±1.77*
Mastitic			7th	$8.76 \pm 0.48$	10.93±0.61	$27.98 \pm 0.89$	14.06±0.68	8.94±0.53	$4.49 \pm 0.42$	4.45±0.34	$1.01 \pm 0.25$	46.13±1.92	26.86±1.48	38.16±1.28
she camels	ment		15th	9.66±0.33	11.18±0.47	$28.98 \pm 0.86$	13.22±0.89	8.44±0.49	4.56±0.27	3.88±0.28	$1.02\pm0.24$	45.75±1.48	24.02±1.39	37.08±1.69
	reat	Gentamicin +	1st	7.30±0.20*	8.05±0.69*	26.20±0.89*	15.83±0.33*	9.03±0.25	$4.20 \pm 0.46$	4.84±0.70*	$0.87 \pm 0.21$	46.86±1.11	26.12±1.65	41.85±1.33*
	post 1	soflupredone acetate	7th	8.59±0.72	11.21±0.51	28.79±0.83	13.59±0.38	$8.44 \pm 0.49$	4.27±0.31	4.17±0.53	1.02±0.19	44.91±1.41	25.39±1.52	37.05±1.55
	щ		15th	9.44±0.82	11.43±0.85	29.30±0.75	13.12±0.94	8.37±0.80	4.38±0.72	3.99±0.65	1.10±0.25	44.10±1.83	23.30±1.70	36.11±1.94

Table 4: Blood picture, Protein profile and Liver enzymes activities of healthy and mastitic she camels

Means with different superscripts within the same column are significantly different at p≤0.05

			-				macro	elements				trace elements						
		Parameters		Serum						nilk		5	Serum (µg/dl	l)		Milk (µg/dl)		ži k _
Groups				Ca mg %	Ph mg %	Na (mEq/L	K (mEq/L)	Ca (mg %)	Ph (mg %)	Na (mEq/L)	K (mEq/L	Copper	Zinc	Iron	Copper	Zinc	Iron	Milk production
		Non		9.38	6.06	136.28	5.23	89.58±	45.64	32.42	65.09	104.52	77.15	82.48	80.63	36.15	178.57	3.95
she camels	tı	reated (control	1)	±	±	±	±	0.75	±	±	±	±	<u>+</u>	±	±	±	±	±
			,	0.73	0.59	0.96	0.44		0.83	0.89	0.87	1.49	1.05	1.23	1.96	1.21	1.52	0.5
		Gentamic	cin	6.09	4.12	133.16	4.82	86.93±	43.95	29.06±	63.24	102.32	76.86	82.10	80.41	35.47±	176.06	4.0
	-			±	±	±	±	0.82	±	0.95	±	±	±	±	±	1.42	±	±
	Itec				0.43*	0.80*	0.63		0.99		0.59	1.60	1.18	1.53	1.75		1.94	0.4
Healthy sl Treated	re	Isoflupred	one	$11.05 \pm$	8.12	135.05	4.07	87.21±	44.28	30.41	63.12	103.11	77.02	81.09	80.34	35.62	176.90	3.9
	Г	acetate		0.38*	±	±	±	0.89	±	±	±	±	±	±	±	±	±	±
					0.48*	0.54	0.20*		0.75	0.64	0.49	1.45	1.09	1.26	1.21	1.14	1.87	0.3
		pre treatment		7.54±	$4.08 \pm$	133.10	4.42±	$86.32 \pm$	43.20	33.06	63.21	102.61	72.50	78.10	78.99	35.14	175.18	2.4
		1		0.29*	0.82*	±	0.74	0.98	±	±	±	±	±	±	±	±	±	4
						0.83*			0.89	0.59	0.54	1.76	1.2*	1.40*	1.84	1.60	1.83	0.3
			1st	7.97	4.49	133.99	4.79	$88.20\pm$	43.98±	32.69	63.59±	103.34	$75.48 \pm$	79.37	79.02±	35.23	176.23	2.6
				±	±	±	±	0.90	0.74	±	0.73	±	1.74	±	1.89	±	±	0.2
		c		0.41*	0.37*	0.43*	0.52			0.87		0.34		1.61		1.72	1.94	
		Gentamicin	7th	9.51	5.82	133.88	$4.94 \pm$	$88.92 \pm$	44.53	32.68	64.68	104.61	76.29	81.09	$80.48 \pm$	35.81	178.41	3.2
		am		±	±	±	0.48	0.63	±	±	±	±	±	±	1.37	±	±	4
		ent		0.91	0.49	0.74			0.61	0.71	0.66	0.48	1.49	1.82		1.48	1.61	0.1
	t.	Ū	15th	9.47	5.87	135.06	5.20	89.30±	45.32	32.69	65.12	104.39	76.99	82.05	80.69	36.08	178.50	3.0
	treatment			±	±	$\pm 0.73$	±	0.51	±	±	±	±	±	±	±	±	±	1
	atn			0.50	0.52		0.55		0.59	0.95	0.39	0.76	1.29	1.58	1.73	1.62	1.05	0.1
			1st	8.14	5.73	134.12	4.63±	$88.04 \pm$	44.09	32.53	64.30±	103.12	76.40	80.19	79.02	35.32	177.30	2.5
	post	plus acetate		±	±	±	0.49	0.84	±	±	0.71	±	±	±	±	±	±	=
	pq	olus		0.61	0.21	0.62			0.33	0.59		0.95	1.23	1.59	1.79	1.47	1.38	0.
			7th	9.55	5.79	135.38	4.98	89.12±	44.52	32.46	64.73	103.87	76.90	81.49	80.28	35.95	178.42	3.4
		Gentamicin plus Isoflupredone aceta		±	±	±	±	0.55	±	±	±	±	±	±	±	±	±	1
		rec		.48	0.17	0.80	0.85		0.29	0.48	0.50	0.78	1.47	1.61	1.37	1.33	1.72	0.4
		lup	15th	9.44	5.90	136.32	5.30	89.63±	44.69	32.40	65.15	104.48	77.17	82.39	80.70	35.18	178.63	3.
		Sof		±	±	±	±	0.48	±	±	±	±	±	±	±	±	±	±
		I		0.62	0.55	0.75	0.33		0.41	0.92	0.38	0.48	1.38	1.40	1.16	1.21	1.31	0.3

Table 5: Some serum, milk elements and milk production of healthy and mastitic she camels

Means with different superscripts within the same column are significantly different at  $p \le 0.0$ 

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#### الملخص العربي

محاولات حقليه لعلاج التهاب الضرع في النوق بمحافظة الشرقية عصام النبراوى ( ايهاب محد حسين \ السيد منصور \ ماجده نعمت عبد الحميد \ رضا حسن ذكى \ فقسم صحة الاغذية - معهد بحوث صحة الحيوان-الزقازيق قسم الكيمياء- معهد بحوث صحة الحيوان-الزقازيق قسم البكتريولوجيا -معهد بحوث صحة الحيوان-الزقازيق

بالفحص البكتيريولوجي لعدد ٣٠ عينة لبن من نوق مصابه بالتهاب الضرع تم عزل البكتيريا المسببة لالتهاب الضرع في النوق ووجد ان ١٨ عينة في صوره عدوى منفردة بنسبه ٢٠% و ١٢ عينة في صوره عدوى مشتركة بنسبه ٤٠%: وبعمل أختبار الحساسية لهذه المعزولات وجد أن الجنتاميسين اكثر المضادات الحيوية تاثيراً على هذه المعزولات. تم إجراء هذا البحث على ٢٥ ناقه بأماكن مختلفه بمحافظة الشرقية (١٥ ناقه سليمه ظاهريا- ١٠ ناقه مصابة بالتهاب الضرع). قسمت النوق إلى ٥ مجموعات متساويه ( ٥ ناقه/ مجموعة) تم حقنها بالجنتاميسين منفردا او مع خلات الأيز وفلوبريدون. تم تجميع عينة لبن من النوق المصابه قبل العلاج وعند ٢, ٢ ١٠ يوم بعد نهاية العلاج لدراسة تأثير التهاب الضرع على كميه وبعض مكونات اللبن. تم تجميع ٢ عينة دم من كل ناقه عند ٢ ٧, ١٠ يوم بعد نهاية العلاج لدر اسة تأثير التهاب الصّرع والعلاج على الصورة الدموية البيوكيميائية. لم يتم عزل البكتيريا المسببه لالتهاب الضرع من اللبن عند ٧ يوم من نهاية العلاج. استخدام الجنتامايسين منفردا او مع خلات الأيزوفلوبريدون كان لهم دور فعال في شفاء الاصابه بالتهاب الضىرع في النوق حيث اختفت الأعراض الظاهرية للمرض وكانت نسبه الشفاء ١٠٠% عند اليوم الخامس والثالث على التوالي من العلاج واديا الي عودة صورة الدم والوظائف البيوكيميائيه إلى المستوى الطبيعي في مصل النوق المصابه والمعالجه بعد العلاج ب ١٠ يوم. الجنتاميسين في النوق أحدثت نقص معنوى في العدد الكلي لكرات الدم الحمراء, تركيز الهيموجلوبين, وحجم كرات الدم المرصوصة% و البروتين الكلي الزلال الجلوبيولين, الكالسيوم, الفوسفور الصوديوم بجانب زيادة معنوية في العدد الكلي لكرات الدم البيضاء, الالنين امينوتر انزفير از , الاسبرتيت امينوتر انزفير از, الفوسفاتيز القاعدي ونقص غير معنوى في البوتاسيوم ومعادن اللبن (الكالسيوم, والفوسفور, الصوديوم والبوتاسيوم) والمعادن النادرة في السيرم واللبن (النحاس الزنك والحديد) خلات الأيزوفلوبريدون أحدثت نقص معنوى في والعدد الكلي لكرات الدم الحمراء تركيز الهيموجلوبين وحجم كرات الدم المرصوصة%, البروتين الكلي, الجلوبيولين بجانب زيادة معنوية في العدد الكلي لكرات الدم البيضاء, الالنين امينوترانزفيراز , الاسبرتيت امينوتر انزفير از ألفوسفاتيز القاعدى الكالسيوم, والفوسفور والبوتاسيوم ونقص غير معنوى في الزلال, الصوديوم معادن اللبن (الكالسيوم, والفوسفور, الصوديوم والبوتاسيوم) والمعادن النادرة في اللبن والسيرم ( النحاس, الزنك التهاب الضرع في النوق أدى الى حدوث نقص معنوى في و العدد الكلي لكرات الدم الحمراء, تركيز الهيموجلوبين وحجم كرات الدم المرصوصةو الزلال والنسبة بين الزلال الجلوبيولين, الكالسيوم, والفوسفور, الصوديوم الزنك الحديد وانتاج اللبن ونقص غير معنوى في البوتاسيوم والنحاس بجانب زيادة معنوية في العدد الكلي لكرات الدم البيضاء البروتين الكلي، الجلوبيولين, الالنين امينوتر انز فير از , الاسبرتيت امينوتر انز فير از, الفوسفاتيز القاعدي مصحوبه بنقص غير معنوي في معادن اللبن (الكالسيوم, و الفوسفور, الصوديوم والبوتاسيوم والنحاس الزنك والحديد) نستخلص من تلك الدر اسة ان التهاب الضرع ادى الى حدوث تأثيرات عكسية على صورة الدم وبعض الوظائف البيوكيميائية ويمكن إستعمال الجنتاميسين وخلات الأبزوفلوبريدون بجرعتهم العلاجيه كعلاج للالتهابات الضرع في النوق.