MYCOPLASMAS PREVALENT IN QUAILS AND SOME WILD BIRDS IN EGYPT

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SUMMARY

The total incidence percentage of mycoplasma which isolated from living quails was 5.75% while it was 3.75% in dead ones, although total recovery rate in dead in sheel eggs was 10% and 6% in infertile quail eggs. Also the total recovery rate of mycoplasma isolated from kestrel falcons was 28.75% and it was 30% in Black-shoulder kite. On the other hand, the total incidence percentage of mycoplasma isolated from Moorhen fowls was 12.5% and it was 20% in Doves. Biochemical tests for glucose and arginine were done and the results were recorded. Serological examination was done by growth inhibition and growth precipitation tests which revealed that 23 strains strains were 21 M.gallisepticum, were M.gallinarum, 11 strains were M. columborale, 9 strains were M.gallinaceum, 4 strains were M. pullorum, one strain was M. synoviae, one strain

was M. columbinum, one strain was M. iners and 32 strains were untyped from quails and wild birds. Serodiagnosis by using ELISA test for both sera and egg yolk of quails had no any detectable antibody titres for both M. gallisepticum and M. synoviae. The results of serodiagnosis of samples collected from living quails which were experimentally infected by M. gallisepticum S 6 strain and M. gallisepticum "field strain" under stress factors by both environmental and biological stresses, proved that no detectable antibodies were present in their sera samples indicating that these birds may act as reservoir for such mycoplasma microorganisms.

INTRODUCTION

During the last two decades a great attention was payed toward quails farming as a trial to fulfill ex-

cessive demands of the increased population from animal protein. Falcons and other wild birds play an important role in environmental balance and such birds are affected by mycoplasma which considered as one of the common poultry pathogens. Bencina et al. (1987) isolated mycoplasma species from six avian species (chicken, chick embryos, turkeys, ducks, geese, pigeons and Japanese quail and their embryos) as 411 out of 792 (52%) were infected with mycoplasmas. Mycoplasma gallisepticum and M. synoviae were the most frequently isolated species, M. anatis was found only in ducks and geese while M. columbinasal, M. columbinum and M. columborale were only present in pigeons. Mycoplasma meleagridis and M. gallopavonis were isolated from turkeys and M. synoviae from pigeons and Japanese qualis. Also different species of mycoplasma were isolated from different poultry species and their eggs and embryos (El-Ebeedy et al., 1987, Kardel, 1987 and Molokwu et al., 1987).

Poveda (1988) found the prevalent mycoplasma species in Peregrino falcon in captivity fed on chicken carcasses, were M. gallisepticum, M. gallinarum, M. gallinaceum and M. iners. The presence of mycoplasma species in wild birds were investigated by some authors as El-Shater et al. (1990), Poveda et al. (1990), Cobb et al. (1992) and Cookson and Shivaprasad (1994). The serodiagnosis of mycoplasma species were done by Optiz et al. (1983), Lin and kleven (1984), Avakian et al. (1988), Dingfelder et al. (1991), Ortiz and

Kleven (1992) and Samuel et al. (1996). This study was planned as an attempt to throw spot lights upon the mycoplasma prevalent in quails and some wild birds via studying the pathogenicity of the prevalent isolated strains and serodiagnosis trial by using enzyme-linked immunosorbent assay (ELISA) for determining the antibody titre for the most pathogenic mycoplasma.

MATERIAL AND METHODS

Samples:

a) Birds: A total of 1020 bacteriological swabs were collected from trachea, lungs, air sacs and genital organs "testicles or ovaries". Including 100 living quails (400 samples), 100 dead quails (400), 20 kestrel falcons (Falco-tinnunculus, 80 samples), 10 Black-shoulder kite (Elanus savigny, 40); 20 Moorhen fowls (Gallinula chloropus, 80) and 5 doves (Columbiformes, 20). In addition to 300 living quails were also used for pathogenicity test.

b) Eggs:

1-<u>Infertile egg</u>s: 100 samples were randomly collected from the egg yolk then suspended in 5 ml PPLO broth for the possible mycoplasmal isolartion and 90 samples out of them were also taken for serological examination.

2- <u>Dead-in shell embryos</u>: 100 samples were taken and then suspended in 5 ml PPLO broth in a trial for mycoplasma isolation.

The basal PPLO broth (Difco) media was prepared and enrichment with the following ingredients: yeast extract 5% Difco (10 ml/100ml), sterile horse serum (15 ml/100 ml), D.N.A. and inhibitors as (2%) thallium acetate (5 ml/100ml) and penicillin G-Na (2000 IU) were aseptically added. The same previously ingredients were also used for preparation of PPLO - agar plates.

Preparation of samples for mycoplasma isolation: was done according to Razin and Tully (1983). Samples were cutted by sterile scissors and grinded with sterile sand, and 5 ml of broth was then added. About 0.2-0.3 ml of the mixture was transferred into the PPLO broth and incubated at 37∞C for 3 days and then subcultured onto PPLO-agar plates which were incubated at 37∞C under reduced oxygen tension in humidified candle jars. Suspected colonies were examined after 48 hours under a stereomicroscope daily up to 7-10 days. The purification and characterization of mycoplasma isolates were done according to Sabry (1968).

Digitonin sensitivity test (Erno and Stipkovitis, 1973) was used for differentiation of myocplasma and acholeplasma colonies. Suspected typical mycoplasma colonies were biochemically identified by using glucose fermentation test and arginine deamination test (Sabry, 1968). Serological identification were done by using growth inhibition test (Clyde, 1964) and growth precipitation test (Krogsgaard, 1972).

Egg yolk preparation for ELISA test (Piela et al., 1985): 0.5 ml from egg yolk examined was taken and inserted in a test tube containing 0.5 ml buffer solution, and 1 ml chloroform solution was added and then centrifuged at 3000 r.p.m. for 3 minutes. The supernatant fluid was used for ELISA test.

Preparation of the serum dilution plate for ELISA test was done according to Mohammed et al. (1985).

ELISA test procedure: Mycoplasma coated test plate must be removed from the protective bag, then 50 ml dilution buffer was added to all wells, then 50 ml diluted mycoplasma positive control serum into well A1, A3 and H11. After that 50 ml/well from the dilution uncoated plate must be transferred to corresponding well of coated testplate. Incubation of such plates was done for 30 minutes at room temperature, then washing three to five times by using automatic washer, then 100 ml diluted conjugate into each well was dispensed. Incubate for 30 minutes at room temperature, then washed again by automatic washer and dispensed 100 ml of substrate solution into each well. Incubated for 15 minutes at room temperture and dispensed 100 ml of stopping solution into each well to stop the reaction. Then measured and recorded absorbance values "optical denisities "O.D." at 405 nm for samples and controls.

The calculation was done by using manual and computerized methods according to Briggs and Skeels (1984).

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Pathogenicity test for Mycoplasma gallisepticum: Preparation of inocula: Local and standard isolates of M.G. were inoculated in PPLO broth without thallium acetate, incubated aerobically at 37∞C for 48 hours and diluted to give an inoculum which contained 105 to 106 C.F.U./ ml. Three hundred quails were divided into 3 groups (each of 100). Each group was subdivided into 5 subgroups (20 quails each). The 1st subgroup were infected intratracheally (I/T) with M. gallisepticum (S6). The 2nd subgroup were intranasally infected (I/N) by M. gallisepticum (S6), while the birds of 3rd ones were intratracheally infected by M. gallisepticum "Field strain". The 4th subgroup were experimentally infected intranasally by M. gallisepticum "field strain", while the birds 5th group was kept as a control. All groups were kept separately. The birds of 1st group received a balanced daily ration and kept in good ventillation condition with good source of light, the birds of 2nd group were stressed by unbalanced ration (containing no adequate amount of both protein and vitamin necessary for their growth) and very bad ventilation condition, while the birds of the 3rd group were biologically stressed by their "La Sota N.D.V. Vaccination" beside unbalanced ration and bad ventilation. The quails of these groups were daily checked to observe any characteristic signs of the C.R.D. syndrome and gross pathological lesions, then 5 birds of these subgroups were weekly slaughtered for post-mortem examination and for mycoplasma reisolation. In addition, sera from birds of the severely stressed group "third group" were collected and serologically examined by ELISA test to investigate the presence of any detectable antibodies.

RESULTS

The recovery rate of mycoplasmas from examined samples were recorded in table (1). The obtained isolates of mycoplasmas were subdivided into three groups according to their biochemical reactions (glucose fermentation and arginine utilization) as shown table (2). Serological typing of such isolates by using growth inhibition less (G.I.T.) and growth precipitation test (G.P.T.) was recorded in table (3) which revealed that from 23 isolates obtained from living quails, 4 strains were related to each of M. gallisepticum and M. pullorum (1% each), 2 strains belonged to M. gallinaceum (0.5%), 6 strains were related to M. gallinaurm (1.5%) and 7 still as untypable strains (1.75%). Meanwhile, from 15 isolates collected from dead quails, 4 strains were related to M. gallisepticum (1%), two strains belonged to M. gallinaceum (0.5%), 3 strains were M. gallinarum (0.75%), one strain belonged to M. iners (0.25%) and five strains were untypable (1.25%). From 15 isolates obtained from quail eggs, 9 strains were related to M. gallisepticum (3 strains originaled from infertile eggs 3% and 6 strains were from dead in shell eggs 6%) and 7 were untypable strains.

As regards to kestrel falcons, M. gallisepticum.

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		rı	sajijese	Al-Azhar fac. of agriculture	Bilbias	Kaír El-Sheikh	Mina Al-Kameh	Abo-Al Matameer	Доліа іе	Total
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	1		No. of birds	30	40	1.1	30	: 35	1 1	100
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			sguni		1	1 1	- 13	: -	1.1	ww
		1	kir sacs			1 1	12 63	1 -	1.1	0.4
		1	genital organe			1.1	2	1 ,	1.1	4 w
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lis.			80	334	6.7 3.13	11	5	2.8	1 1	5.75 3.75
	T	_	No. of cases	1 20	20	11	: 6	20	: :	: 8
1.8	ı	Infertile	Myco. (+)	1 1	110	1 1	; ω	1 -	1 1	: 0
E.	١	7	ઋ	1 1	- 10	1 1	7.5	1 00	11	: 6
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		SS33	89	5!	6:	v i	125	1 1	1.1	5:
to A	_		No. of birds	11	0:	1.4	1 1	1:	10 :	20
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	Slwc	345	sacs niA	1 1	10 1	- 1	11	11	1 1	w ;
			genital organe	1 1	11	11	1.1	11	1 1	1.1
			89	: 1	16.7	:- 125	11	11	10 1	12.60
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	Falcons	160	saunt	1 1	1.1	2	11	11	1.1	13 -
	₩.	5	Socs niA	11	– ω	- 6	1 1	1.1	1.1	12 9
			genital organe	1 1	11	1.1	1.1	1.1	1 1	11
		Anna.	%	1 1	25 50	31.25 27.80	1 1	1.1	1.1	28.75 30
	L	aqYT	and the state of the state of	1 1	w ×	m ×	1.1	1.1	11	BX
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		8	genital organe	11	1 1	1-1	1 1	11	1.1	11
			%	1.1	20	1 1	1.1	1 1	1.1	20

Table 1: Recovery rate of mycoplasma from quails and some wild birds.

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Table 2: Biochemical characterization of mycoplasma species isolated from quails and some wild birds.

	Criteria	1	Site of isolation	No of Samples*	Gl. (+) Ar. (-)	Gl. (-) Ar. (+)	Gl. (+) Ar. (+)
	Birds	Living Quails	Trachea Lungs Air-Sacs Genital	8 5 6 4	5 3 3 4	2 2 3 	1 :: ::
\overline{Q}	Sp		Sum	23	15	7	1
ds Quails		Dead Quails	Trachea Lungs Air-Sacs Genital	5 3 4 3	3 1 3 3	2 2 1 	
			Sum	15	10	5	- 1
	Eggs	Infertile Dead-in-shell		6 10	6 9	 1	
	Sg	Dead-III-sticii	Sum	16	15	Ī	-
H	Kestrel F.		Trachea Lungs Air-Sacs Genital	13 9 1	8 1 2 	5 1 	
2	2		Sum	23	11	6	
COARD	Kalcons	Blackshoulder kite F.	Trachea Lungs Air-Sacs Genital	8 2 2 	6 1 2 	2 1 	
		feets	Sum	12	9	3	d and see
	Moorhen	Fowls	Trachea Lungs Air-Sacs Genital	5 2 3 	4 2 2 	1 1 	
		4	Sum	10	8	2	
Doves		Trachea Lungs Air-Sacs Genital	3 1	2	1 1 1	- -	
		*_ 1	Sum	4	2	2	
	Tot	al		103	70	26	1

^{*}All samples were sensitive to 1.5% digitonin.

Ar.: Arginine deamination test.

Gl. Glucose fermentation test.

Table (3): Results of serotyping of mycoplasma isolates by "G.LT," according to biochemical grouping.

			,	Qual	sam	ples				alcon s	amole			_		
-1			Bird	s			Egg	s		arcon a		vo		doord	Do	
Type of mycoplasmal isolates	LIVING (400)		Dead (400)		Internie (100)	16.2.1000	Eg	gs		Kestrel F. (80)	klte (40)	Bkack-shoulder	(80)	Moorhen fowls samples	Doves samples (20)	
al isolates	No. of isolates	Incidence %	No. of isolates	Incidence %	No. of isolates	Incidence %	No. of isolates	Incidence %	No. of isolates	Incidence %	No. of isolates	Incidence %	No. of isolates	Incidence %	No. of isolates	Incidence %
Group (1): M. gallisepticum M. synoviae M. pullorum	4	1	4	Î	3	3	6	6	2	2.5	1	2.5	3,	3.75 1.75		
M. givcophilum M. gallinaceum M. anatis M. gallopavonis M. columborale	2	0.5	2	0.5					8	10	1	2.5	4	5	2	10
Untyped	5	1.25	4	1	3	3	3	3	7	8.75	6	15	T di			ulta.
Group (2): M. anseris M. cloacale M. meleagridis M. gallinarum M. iners M. columbinasal	6	1.5	3	0.75 0.25					6	7.5	3	7.5	2	2.5	-	5
M. columbinum Unt yped	1	0.25	1	0.25			1.	1								Ĺ
Group (3): M. iowae M. lipofaiens Untyped	1	0.25									r 182					
Total	23	5.75	15	3.75	6	6	10	10	23	28.75	12	30	10	12.5	4	20

G.I.T. Growth inhibtion test

G.P.T. Growth precipitation test

M, columborale, M. gallinarum, and untypable isolates were identified in an incidence of 2.5 %, 10%; 7.5% and 8.75% respectively.

From 12 isolates obtained from Black-shoulder-kite falcons, M. gallisepticum, M. gallinaceum, M. columborale and M. gallinarum were identified in an incuidence of 2.5%, 2.5%, 23.5% and 7.5% respectively and 6 strains still untypable (15%). From 10 isolates collected from Moorhen fowls, 3 strains belonged to M. gallisepticum (3.75%), one strain belonged to M.synoviae (1.25%), 4 strains were related to M. gallinaceum (5%) and two strains belonged to M. gallinarum (2.5%).

From 4 isolates obtained from doves, 2 strains were related to *M. columborale* (10%), one strain was each of *M. gallinarum* and *M. columbinum* (5% each).

Frequency distribution of mycoplasma isolated from various sites:

A total of 8 isolates were recovered from trachea of living quails (2%), 2 were identified as *M. gallisepticum* (0.5%), one isolate was each of *M. pullorum*, *M. gallinaceum*, *M. gallinarum* (0.25% each) and 3 isolates were untypable (0.75%). Five isolates were obtained from lungs of living quails (1.25%), two of them were identified as *M. pullorum* (0.5%), one isolate was *M. gallinaceum* (0.25%) and two isolates were *M. gallinarum*

(0.5%). Six isolates were isolated from air sacs of living quails (1.5%), one isolate from each of M, gallisepticum and M, pullorum (0.25% each), 3 isolates were M, gallinarum (0.75%) and one isolate was untypable (0.25%).

Four isolates were recovered from genital organs of living quails (1%) "2 isolates from the ovaries and 2 isolates from testes", one isolate was M. gallisepticum (0.25%) and the other 3 isolates were untypable (0.75%). In addition to five isolates were obtained from trachea of dead quails (1.25%) from both one isolate from each of M. gallisepticum, M. gallinaceum, and M. iners (0.25% each) was identified and two isolates were untypable (0.5%). Also, three isolates were collected from lungs of dead quails (0.75%), one of them was M. gallinaceum (0.25%) and two of them were indentified as M. gallinarum (0.5%).

Four isolates were secured from air sacs of dead quails (1%), one isolate from each of *M. gallisepticum* and *M.gallinarum* (0.25% each) and the remaining were untypable (0.5%). Lastly, three isolates were obtained from quail genital organs (0.75%), two of them were identified as *M. gallisepticum* (0.5%) "one isolate from ovary and another one from testes" and one isolate still untyped was recovered from ovary. Sixteen isolates were recovered from quail eggs, six isolates were from infertile eggs (6%), three of them were serologically typed as *M. gallisepticum* (3%) and 10 isolates were from dead in shell eggs (10%), six

of them were M. gallisepticum (6%) and 7 isolates were untypable (7%) as shown in table (4).

Thirteen isolates were collected from the trachea of Kestrel falcons (16.25%), two of them were identified as M. columborale (2.5%), one isolate was M. gallisepticum (1.25%), five of them belonged to M. gallinarum (6.25%) and the other five isolates were untypable (6.25%). Also one untyped isolate was obtained from lung of kestrel falcons (1.25%). While nine isolates were collected from air sacs of kestrel falcons (11.25%), six of them were M. columborale (7.5%), one isolate was identified as each of M. gallisepticum, M. gallinarum and untyped strain (1.25% each). Eight isolates were collected from trachea of Black-shoulder-kite falcons (20%), on identification, one of them was M. gallisepticum (2.5%), two isolates were M. gallinarum (5%), one isolate was M. gallinaceum (2.5%) and four isolates were untypable (10%). Also two isolates were secured from lungs of Black-shoulder-kite falcons (5%), one of them was M. gallinaurm (2.5%) and another one was still untyped (2.5%). Besides two isolates were obtained from air sacs of Black shoulder-kite falcons (5%), one isolate was M. columborale (2.5%) and one isolate was untyped (2.5%). No isolates could be obtained from genital organs of falcons as shown in table (4).

Five isolates were collected from trachea of Moorhen fowls (6.25%), one of them was each of M. gallisepticum, M. gallinarum and M. synoviae (1.25% each) and two isolates were identified as

M. gallinaceum (2.5%). Also two isolates were recovered from lungs of Moorhen fowls (2.5%), and identified as M. gallisepticum and M. gallinaceum (1.25% each). While three isolates were obtained from air sacs of Moorhen fowls (3.75%), on identification, M. gallisepticum, M. gallinaceum and M. gallinarum were recorded (1.25% each).

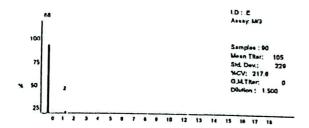
Three isolates were obtained from trachea of doves (15%), two isolates belonged to *M. columborale* (10%) and one was *M. gallinarum* (5%). While only one strain belonging to, *M. columbinum* (5%) was obtained from air sac. No isolates could be obtained from lungs and genital organs of doves. The previously data are recorded in table (4).

Results of serodiagnosis of *M. gallisepticum* and *M. synoviae* by using ELISA technique:

A) For *M. gallisepticum*: The results of serodiagnosis of *M. gallisepticum* revealed that only two serum samples gave positive immunoresponse at 1st group of titre (2.22%) and the remaining 88 serum samples lied in (0) group of titre (97.8%) and the mean titre was 105 under the mean titre of group "0" (0-499) as in (Fig.1). While in quail eggs the obtained data indicated that two sera out of examined 90 samples were positive at 1st group of titre (2.22%) and one sample was positive each of 2nd and 3rd titre group (1.11%) and (1.11%)

(80.	3	kite (40)	(80 fa 7	`	(400)	(400) C	N.
Doves	Moorhen Fowl (80 samples)	kite falcons (40 samples)	falcons (80 samples)		ails ples	Living Quails (400) samples	of samples and isolations
Trachea Lungs Air-Sacs	Trachea Lungs Air-Sacs Total (80)	Trachea Lungs Air-Sacs Total (80)	Trachea Lungs Air-Sacs Total (80)	Inferile (100) Dead inshell (100) Total	Trachea Lungs Air-Sacs Genital ovary organs testes Total (400)	Trachea Lungs Air-Sacs Genital ovary organs testes Total (400)	d site of
	3	1	1 - 1 2) 6 9	4-1	42	gallisepticum No.of isolates
	1.25 1.25 1.25 3.75	2.5 2.5	1.25 1.25 2.5	9	0.25 0.25 0.25 0.25 1	0.5 0.25 0.25 1	%
	, , , ,			No.		4 2 -	M. pullorum No.of isolates
	1	11.1				0.25 0.5 0.25	%
	44				21.122	2111-	M. gallinaceum No of % isolates %
	2.5 1.25 1.25 5	2.5 - 2.5			0.25 0.25	0.25 0.25 	and the sale
	1	2 1 3	6-1.5		ωυ·	611011	galinarum No.of isolates
۸ ، ، ۸	1.25 1.25 2.5	5 2.5 7.5	6.25 - 1.25 7.5		0.5 0.25	0.25 0.5 0.75 -	% m
				. 77	211112		iners No.of
					0.25		%
21112			8 O · 12				M. columborale
5 · · 6	eale da	5.5	2.5 7.5 10				
			* 1 1 1				M. synoviae
Market A	1.25 1.25						% ac
							M. columbinum No.of %
					5 A 3 C C 1		%
		40		υ4 Γ	v - 12 - 12	7-23	Untypable mycoplasma No.of % isolates
		10 2.5 2.5	6.25 1.25 1.25 8.75	7 43	0.5 0.5 0.25	0.75 0.25 0.25 0.25	able lasma
3	5 2 2 5					222058	Total No.of isolates
15	6.25 2.5 3.75 12.5	50 30	16.25 1.25 11.25 28.75	16	1.25 0.75 1 0.5 0.25	2 1.25 1.5 0.5 0.5 5.75	%

Table(4): Frequency distribution of mycolplasmas isolated from quails and some wild birds.



E MG DIL:1:500 NEAN TITER: 105 % CV: 217.6 G.M.T.: 0

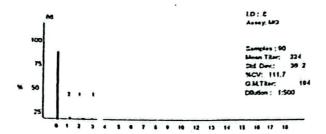
G.M.T		0 .						
WELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP		WELL	RAW O.D.	S/P RATIO
- A01	0.147	00.04	72	0	1	44 D12	0.0158	00.06
- A02	0.093	00.00	0	0		45 E01	0.0198	00.12
+ A03	0.895	00.90	2047	3		46 E02	0.001	00.00
+ A04	0.820	01.10	2538	3		47 E03	0.188	00.11
I A05	0.163	00.07	121	0		48 E04	0.089	00.00
2 A06	0.182	00.10	180	0		49 E05	0.377	00.40
3 A07	0.229	00.17	334	0		50 E06	0.028	00.00
4 A08	0.132	00.02	30	0		51 E07	0.054	00.00
5 A09	0.190	00.11	208	0	1	52 E08	0.073	00.00
6 A10	0.194	00.12	219	0		53 E09	0.035	00.00
7 AII	0.080	00.00	0	0		54 E10	0.045	00.00
8 A12	0.011	00.00	0	0		55 E11	0.091	00.00
9 B01	0.042	00.00	0	0		56 E12	0.026	00.00
10 B02	0.511	00.61	1344	i i		57 F01	0.004	00.00
11 B03	0.267	00.23	462	0 -		58 F02	0.122	00.00
12 B04	0.151	00.05	84	0		59 F03	0.122	00.00
13 B05	0.249	00.20	401	0		60 F04	0.082	00.00
14 B06	0.180	00.09	174	0	1	61 F05	0.176	00.09
15 B07	0.207	00.14	261	0		62 F06	0.047	00.00
16 B08	0.225	00.16	320	0		63 F07	0.031	00.00
17 B09	0.121	00.00	2	0	-	64 F08	0.020	00.00
18 B10	0.041	00.00	0	0		65 F09	0.018	00.00
19 B11	0.042	00.00	0	0	-	66 F10	0.025	00.00
20 B12	0.001	00.00	0	0		67 F11	0.007	00.00
21 C01	0.034	00.00	0	0		68 F12	0.025	00.00
22 C02	0.086	00.00	0	0		69 G01	0.044	00.00
23 C03	0.117	00.00	0	0		70 G02	0.039	00.00
24 C04	0.192	00.11	212	0		71 G03	0.144	00.04
25 C05	0.457	00.53	1143	0		72 G04	0.036	00.00
26 C06	0.127	00.01	16	0		73 G05	0.076	00.00
27 C07	0.127	00.16	310	0	-	74 G06	0.199	00.12
28 C08	0.163	00.07	121	0	1	75 G07	0.033	00.00
29 C09	0.060	00.00	0	0		76 G08	0.299	00.28
30 CIO	0.000	00.00	Ö	0		77 G09	0.058	00.00
31 C11	0.013	00.00	Ö	0		78 G10	0.068	00.00
32 C12	0.022	00.00	Ŏ	Ō	ı	79 G11	0.009	00.00
33 D01	0.022	00.03	49	ŏ	1	80 G12	0.072	00.00
34 D02		00.03	267	ŏ	l	81 H01	0.133	00.02
35 D03	0.209	00.00	0	ŏ	١	82 HO2	0.124	00.01
36 D03	0.085		ő	ő		83 H03	0.094	00.00
37 D05	0.084	00.00	0	ŏ		84 H04	0.097	00.00
38 D06	0.067	00.00	459	ŏ		85 H05	0.062	00.00
39 D07	0.266	00.23		ő		86 H06	0.104	00.00
The second secon	0.110	00.00	0	ŏ	4 (87 H07	0.212	00.14
40 D08 41 D09	0.101	00.00	0	ŏ		88 H08	0.024	00.00
42 D01	0.030	00.00	0	Ö		89 H09		00.00
- 111	0.009	00.00	0	Ö				00.00
43 DII	0.019	00.00	0	U		90 H10	0.046	00.00

Fig. (1): Mycoplasma gallisepticum antibody titre in sera of quails.

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TITR GROUP

Titre



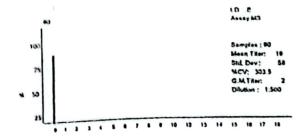
E MG DIL:1:500

	NEAN G.M.T	TITER	: 324 % CV: 194	111.7				
W	ELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP	1	WELL	RAW O.D.
-	A01	0.005	00.00	1	0		44 DI	0.033
-	A02	0.010	00.01	io	ŏ	1 1	45 EO	36010000000000
+	A03	0.418	00.15	2663	3		46 E02	
+	A04	0.312	00.85	1923	2		47 E03	
1	A05	0.083	00.33	420	ő		48 E04	
2	A06	0.019	00.03	54	ŏ		49 E05	
3	A07	0.033	00.07	128	ő	1 1	50 E06	
4	A08	0.029	00.06	106	ŏ	1 1	51 E07	
5	A09	0.069	00.17	336	ŏ	F 1	52 E08	
6	A10	0.384	01.05	2423	3		53 E09	
7	AII	0.075	00.19	372	ő		54 E10	
8	A12	0.193	00.52	1120	ĭ		55 EII	The state of the s
9	B01	0.286	00.78	1744	2		56 E12	
10	B02	0.060	00.15	283	ő			0.142
11	B03	0.035	00.08	139	Ö			0.031
12	B04	0.069	00.17	336	ŏ		58 F02	The second second second
13	B05	0.015	00.02	33	Ö		59 F03	0.028
14	B06	0.020	00.02	59	ŏ		60 F04	0.059
15	B07	0.073	00.03	360	ő		61 F05	0.048
16	B08	0.013	00.02	24	ő		62 F06	0.053
17	B09	0.063	00.02	300	ő		63 F07	0.061
18	B10	0.003	00.10	38	ő	4	54 F08	0.019
19	BII	0.088	00.02	451	ŏ		55 F09	0.213
20	B12	0.023	00.04	74	ő	A	56 F10	0.035
21	COI	0.023	00.07	134	ŏ		57 FII	0.019
22	C02	0.047	00.07	207	ő		58 F12	0.035
23	C03	0.105	00.27	555	ő		59 G01	0.026
24	C04	0.083	00.21	420	ŏ		70 G02	0.163
25	C05	0.051	00.12	230	ŏ		71 G03	0.091
26	C06	0.105	00.12	555	ŏ		2 G04	0.017
27	C07	0.103	00.10	184	ŏ		3 G05	0.071
28	C08	0.144	00.38	802	ŏ		4 G06	0.052
29	C09	0.056	00.14	259	ŏ		5 G07	0.057
30	CIO	0.066	00.16	318	ŏ		6 G08	0.146
31	CII	0.030	00.06	112	ŏ		7 G09	0.088
32	CI2	0.039	00.09	162	ŏ		8 G10	0.097
33	D01	0.030	00.06	112	ŏ		9 GII	0.004
34	D02	0.045	00.10	196	ŏ		0 G12	0.023
35	D03	0.002	00.00	1	ŏ		H01	0.64
36	D04	0.065	00.16	312	ŏ		2 H02	0.133
37	D05	0.015	00.02	33	ŏ		3 H03	0.066
38	D06	0.017	00.03	43	ő		4 H04	0.065
39	D07	0.101	00.26	531	ŏ		5 H05	0.049
40	D08	0.031	00.07	117	ŏ		6 H06	0.086
41	D09	0.042	00.10	179	ŏ		7 H07	0.043
42	DOI	0.027	00.05	96	ŏ		8 H08	0.075
43	DII	0.041	00.09	173	ŏ	0.00		0.077
7.5	-	A MILAN	A STATE OF THE PARTY OF THE PAR	The Same	and the same of	19	0 H10	0.057

Fig. (2): Mycoplasma gallisepticum antibody titre in eggs of quails.

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E MG DIL:1:500 NEAN TITER: 19 % CV: 303.5

	G.M.T.	;	2		
W	/ELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP
	A01	0.098	00.00	1	0
	A02	0.145	00.04	71	0
	A03	0.508	00.68	1507	2 4
+	A04	0.866	01.32	3097	
1	A05	0.014	00.00	_1	0
	A06	0.147	00.05	78	0
3	A07	0.017	00.00	1	0
4	A08	0.083	00.00	1	0
5	A09	0.005	00.00	1	0
6	A10	0.034	00.00	!	0
7	AII	0.055	00.00	1 1	0
8	A12	0.088	00.00	1	0
9	B01	0.139	00.03	51	
10	B02	0.143	00.04	65	0
11	B03	0.129	00.01	20 I	l ŏ l
12	B04	0.072	00.00	74	l ŏ l
13	B05	0.146	00.04		l ŏ l
14	B06	0.170	00.09	157	l ŏ l
15	B07	0.094	00.00	228	l ŏ l
16	B08	0.189	00.12	1 220	ŏ
17	B09	0.082	00.00	172	l ŏ l
18	B10	0.174	00.09	172	l ŏ l
19	B11	0.116	00.00	240	l ŏ l
20		0.193	00.13	277	l ŏ l
21	COL	0.203	00.14	211	ŏ
22	C02	0.071	00.00	ì	l o l
23	C03	0.047	00.00	1	Ö
24		0.032	00.00	l i	0
25		0.026	00.00	i i	0
26	C06	0.029	00.00	l i	0
27		0.030	00.00	l i	0
28		0.044	00.00	ì	0
29		0.065	00.00	ì	0
30		0.054	00.00	Î.	0
31	CII	0.071	00.00	ĺ	0
32	C12	0.020	00.00	1	0
33		0.007	00.00	1	0
34		0.004	00.00	1	0
35		0.013	00.00	1.	0
30		0.013	00.00	1	0
37		0.003	00.00	1	0
38		0.073	00.00	1	0
39		0.001	00.00	1	0
40		0.001	00.00	1	0
4		0.031	00.14	270	0
4		00.084	00.00	1	. 0
4	3 DII	00.00		حسل	W 12 - 1 - 1 - 1

W	ELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP
44	D12	0.006	00.06	4	0
	E01	0.008	00.00	Ī	. 0
45	Total 180	0.088	00.00	ì	0
46	E02	0.093	00.00	î	0
47	E03		00.00	ì	0
48	E04	0.066	00.00	i	0
49	E05	0.013	00.00	. ĝ.	0
50	E06	0.048	00.00	i	0
51	E07	0.005	00.00	î	0
52	E08	0.091	and the second second	i	ŏ
53	E09	0.95	00.00	i	ő
54	E10	0.094	00.00	i	ŏ
55	EII	0.027	00.00		ő
56	E12	0.097	00.00	i	ő
57	F01	0.049	00.00	1	ŏ
58	F02	0.014	00.00	i	ŏ
59	F03	0.012	00.00		ő
60	F04	0.052	00.00		ŏ
61	F05	0.001	00.00	1	ő
62	F06	0.042	00.00		ő
63	F07	0.009	00.00		ŏ
64	F08	0.095	00.00	1	Ĭ
65	F09	0.094	00.00	1	0
66	F10	0.075	00.00	ļ	Table 2 co
67	F11	0.079	00.00	!	0
68	F12	0.049	00.00	ı İ	0
69	G01	0.035	00.00		0
70	G02	0.041	00.00	1	0
71	G03	0.022	00.00	1.	0
72	G04	0.081	00.00	1	0
73	G05	0.080	00.00		0
74	G06	0.105	00,00		0 -
75	G07	0.062	00.00	l l	0
76	G08	0.03	00.00		0
77	G09	0.039	00.00	1	0
78	G10	0.052	00.00	1	0
79	GII	0.058	00.00		0
80	GI2	0.091	00.00	1	0
81	H01	0.048	00.00	. 1	0
82	H02	0.038	00.00	1	0
83	H03	0.018	00.00	1	0
84	H04	0.061	00.00	1	0
85	H05	0.058	00.00	1	0
35.000	H06	0.037	00.00	1	0
86	H07	0.090	00.00	l i -	0
87	H08	0.083	00.00	Î	0
88	F 3 E 5	0.033	00.00	i	0
89	H09	0.113	00.00	l i	0
90	H10	0.003	00.00	200	the state of

Fig. (3): Mycoplasma synoviae antibody titre in sera of quails.

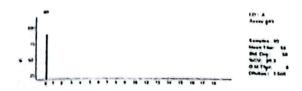
E MG DIL:1:500 NEAN TITER: 79 % CV: 206.8

G.M.T.:		16		
WELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP
WELL - A01 - A02 + A03 + A04 1 A05 2 A06 3 A07 4 A08 5 A09 6 A10 7 A11 8 A12 9 B01 10 B02 11 B03 12 B04 13 B05 14 B06 15 B07 16 B08 17 B09 18 B10 19 B11 20 B12 21 C01 22 C02 23 C03 24 C04 25 C05 26 C06 27 C07 28 C08 29 C09 30 C10 31 C11	O.D. 0.004 0.008 0.315 0.225 0.011 0.006 0.001 0.002 0.038 0.007 0.016 0.027 0.012 0.008 0.002 0.001 0.005 0.017 0.003 0.002 0.004 0.023 0.004 0.023 0.004 0.023 0.004 0.010 0.006 0.010 0.006 0.011 0.015 0.018 0.025	S/P RATIO 00.00 00.01 00.17 00.83 00.02 00.00 00.00 00.00 00.01 00.02 00.01 00.02 00.01 00.00 00.04 00.08 00.02 00.01 00.00 00.00 00.00 00.00 00.00 00.06 00.00 00.06 00.00 00.06 00.00 00.06 00.00 00.01 00.05 00.00 00.00	2 11 2719 1868 30 2 2 2 2 22 22 2 2 2 107 2 115 2 107 2 115 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	
30 C10	0.018	00.05 00.07 00.12 00.04 00.34 00.03 00.03 00.00 00.00 00.11 00.03 00.03	78 130 221 71 708 50 50 2 2 206 57 43	0 0 0 0 0 0 0 0
42 D01 43 D11	0.013 0.016	00.03 00.04	43 64	0

W	ELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP
11	D12	0.032	00.10	183	0
44	100		00.00	2	0
45	E01 E02	0.004 0.006	00.00	2	0
46 47	E02	0.000	00.00	17	0
47	E03	0.009	00.01	50	0
49	E05	0.003	00.00	2	0
50	E06	0.003	00.00	2	0
51	E07	0.004	00.00	17	0
	E08	0.009	00.01	43	0
52 53	E09	0.013	00.05	100	0
53 54	E10		00.07	122	0
	EII	0.024	00.07	167	0
55 56	E12	0.030 0.033	00.09	190	0
57	F01	7.7.7.7.7.7	00.00	2	0
58	F02	0.006 0.022	00.06	107	0
59	F03	0.022	00.00	2	Ö
60	F04		00.00	23	0
61	F05	0.010	00.02	2	0
62	F06		00.00	30	0
63	F07	0.011	00.02	30	ő
64	F08	0.011	00.02	137	0
65	F09	0.026	00.08	198	Ö
66	F10	0.034 0.043	00.11	269	0
67	FII		Approximation of the second	308	O
68	F12	0.048	00.16	333	0
69	G01	0.051	00.17 00.00	5	l ő
70	G02	0.007		2	0
71	G02	0.003	00.00	30	0
72	G04	0.011	00.02 00.01	17	0
73	G05	0.009			0
74	G06	0.001	00.00	2 2 5	0
75	G07	0.005	00.00	2	0
76	G08	0.007	00.00	2	0
77	G09	0.000		37	0
78	G10		00.02	3/	0
79	GII	0.014	00.03	50	0
80		0.004	00.00	2	0
81	H01	0.007	00.00	5	Decidence of the Control of the Cont
82		0.013	00.03	43	0
		0.017	00.04	71	0
83	10 P.	0.007	00.00	5 2 2 2 2 2 2	0
84	7 7 7 7	0.001	00.00	2	0
85		0.006	00.00	2	0
86	F 97000 300	0.003	00.00	2	0
87		0.004	00.00	2	0
88		0.005	00.00		0
89		0.008	00.01	.11	0
90	H10	00.09	00.01	17	0

Fig. (4): The level of Mycoplasma synoviae in quail eggs.

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E MG DIL:1:500 NEAN TITER: 58 % CV: 96.3 G.M.T.: 0

G.M.T.	:	0		
WELL	RAW O.D.	S/P RATIO	Titre	TITR GROUP
- A01	0.004	00.00	0	0
- A02	0.004	00.00	0	0
+ A03	1.294	01.00	2287	3
+ A04	1.297	01.00	2293	3
1 A05	0.030	00.02	32	0
	0.021	00.01	20	0
2 A06 3 A07 4 A08	0.006	00.00	1	0
4 A08	0.043	00.03	50	0
5 A09	0.050	00.04	60	0
6 A10	0.047	00.03	58	0
7 All	0.065	00.050	82	0
8 A12	0.035	0.02	39	0
9 B01	0.025	00.02	25	0
10 B02	0.073	00.05	94	0
11 B03	0.071	00.05	91	0
12 B04	0.011	00.01	7	0
13 B05	0.121	00.09	167	0
14 B06	0.057	00.04	70	0
15 B07	0.035	00.02	39	0
16 B08	0.079	00.06	102	Ö
Service and the service and th	0.025	00.02	25	Ö
	0.52	00.02	63	ő
	0.005	00.00	0	Õ
19 B11		00.03	58	ŏ
20 B12	0.047	00.03	28	Ö
21 C01		00.02	19	Õ
22 C02	0.020	00.01	39	ő
23 C03	0.035	00.02	39	ő
24 C04	0.035		39	ŏ
25 C05	0.035	00.02	0	ő
26 C06	0.005	00.00	6	ŏ
27 C07	0.010	00.00	6	Ö
28 C08	0.010	00.00	63	ő
29 C09	0.052	00.04	33	ŏ
30 C10	0.031	00.02	25	ő
31 CII	0.025	00.02		ő
32 C12	0.019	00.01	17 133	ő
33 D01	0.099	00.07		Ö
34 D02	0.062	00.04	77	Ö
35 D03	0.091	00.07	121	365
36 D04	0.010	00.00	6	0
37 D05	0.036	00.02	40	0
38 D06	0.016	00.01	13	
39 D07	0.033	00.02	36	0
40 D08	0.111	00.08	151	0
41 D09	0.016	10.00	13	0
42 DOI	0.050	00.04	60	0
43 DII	0.046	00.03	54	0

WELL		RAW O.D.	S/P RATIO	Titre	TITR GROUP
44	D12	0.003	00.00	0	0
45	EO!	0.030	00.02	32	0
46	E02	0.115	00.09	157	0
47	E03	0.137	00.10	192	0
48	E04	0.213	00.16	314	0
49	E05	0.045	00.03	53	0
50	E06	0.034	00.02	37	0
51	E07	0.032	00.02	35	0
52	E08	0.105	00.08	142	0
53	E09	0.028	00.02	29	0
54	E10	0.09	00.08	148	0
55	E11	0.062	00.04	77	0
56	EI2	0.013	00.01	10	0
57	F01	0.038	00.03	43	0
58	F02	0.058	00.04	71	0
59	F03	0.069	00.05	88	0
60	F04	0.054	00.04	66	0
61	F05	0.070	00.05	89	0
62	F06	0.001	00.00	0	0
63	F07	0.025	00.02	25	0
64	F08	0.023	00.02	33	0
65	F09	0.045	00.03	53	ī
66	F10	0.084	00.06	110	0
67	FIL	0.023	10.00	23	0
68	FI2	0.028	00.02	29	0
69	GOI	0.125	00.02	173	Ö
70	G02	0.012	00.01	8	0
71	G03	0.012	00.01	7	Ö
72	G04	0.072	00.05	92	Ö
73	G05	0.072	00.02	32	ő
74	G06	0.014	00.02	11	ŏ
75	G07	0.014	00.01	16	Ö
76	G08	0.114	00.01	156	ő
77	G09	0.026	00.02	29	Ö
78	G10	0.020	00.02	36	ŏ
	GII	A	00.02	27	ő
79	GI2	0.026	00.02	28	0
80	The state of the s	0.027	The second second second	225	ő
81	H01	0.158	00.12	66	0
82	H02	0.054	00.04	11	0
83	H03	0.014	00.01	59	0
84	H04	0.049	00.03		0
85	H05	0.107	00.08	145	0
86	H06	0.039	00.03	44	10.00
87	H07	0.046	00.03	54	0
88	H08	0.052	00.04	63	0
89	H09	0.006	00.00	1	0
90	H10	0.011	00.01	7	0

Fig. (5): Level of Mycoplasma gallisepticum antibodies in sera of group of quails experimentally infected with this microorganism under stress conditions.

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for each respectively, while the remaining 86 samples gave no detectable immunoresponse (95.6%) and the mean titre was 324 under the mean titre group "O' (0-499) as shown in (Fig. 2).

B) For *M. synoviae*: The result of the serodiagnosis of *M. synoviae* by using ELISA technique revealed that all collected isolates obtained from sera of the examined 90 quails were presented under titre group (0) 100% as specific antibodies did not present and the mean titre were (19), Geometric mean titre were (2), which lies under group zero (0-499) as illus-

trated in (Fig. 3).

The *M. synoviae* antibodies titre in quail eggs revealed that the one smaple was lied under the group (I) "1.11%" and the remaining 89 samples gave no detectable immunoresponse (98.9%). The mean titre was (79), Geometric mean was (16) which lies under group Zero (0-499) as in (Fig.4).

Results of experimental infection of quails with both local field strain and S6 strain of *M. gallisepticum*:

The obtained data revealed that all groups of quails bred under various stress factors did not clinically succumb the infection and there were no characteristic *P.M. lesions* of *M. gallise*pticum infection. When serum samples of quails of the

3rd group "stressed by both environmental and bi. ological stresses" were collected and serologically examined by ELISA they gave no detectable specific antibodies (Fig. 5).

DISCUSSION

Like most creatures quails and wild birds may play a considerable role in the dissemination of many pathogens and their role for the transmis. sion of mycoplasma organisms in still a point of argument, so this study was planned to detect their anticipated role as reservoirs for these microorganisms. In order to attain this goal random samples were collected from quails, Falcons (Kcstrel and Black-shoulder kite), Moorhen fowls and Doves. The obtained data revealed that 23 mycoplasma isolates were recovered from 400 samples collected from 100 living quails (5.75%). The serological identification and frequency distribution of such isolates were recorded in tables (3&4) Six mycoplasma isolates originated from 100 in fertile quail eggs (6%), three out of them wer M.gallisepticum (3%) and 3 were untyped (3%) In addition to 10 mycoplasma isolates were so cured from 100 dead in shell embryos egg (10%), 6 out of them were M. gallisepticum (65) and 4 were untyped (4%). These results agree with whose reported by Tiong (1978) who isola ed and identified M. gallisepticum from quails, suffering from contagious purulent simb itis. Moreover, Maria and Elmiro (1985) isolale



M. gallisepticum from sinuses of Japanese quails. Also, Reece et al. (1986A) isolated M. gallisepticum from two flock of quails but with high incidence rates of 10% and 13%. While Shah-Majid and Nihayah (1987) proved that no mycoplasmas could be isolated from 20 examined quails indicating that they did not contracted the infection. Also, Cookson and Shivaprasad (1994) detected mycoplasmal from samples collected from quails and added that such isolates were other than M. gallisepticum, M. synoviae, M. meleagridis or M. iowae. On the other hand, there was a gap between the obtained findings and that repoted by Lin and Liu (1985) who found that the mycoplasmas were isolated from tracheal swabs of 209 samples out of 437 fowls, M. gallisepticum (27%), M. synoviae (42%), M. gallinarum (56%), (9%) and M. gallipavonis M. gallinaceum (8.6%), were identified and near these incidence rates were obtained by Kardel (1987). These may be attributed to the difference in environment of quails.

As regards to the incidence of mycoplasmas in some wild birds, the obtained results were supported by the findings of Howse and Jordan (1983), Poveda (1988) and Poveda et al. (1990) who found 13 mycoplasma isolates in various types of flacons. Six isolates could be identified, 3 isolates from black vulture were *M. gallinarum*, 2 isolates from peregrine falcons were *M. columborale* and one isolate from suker falcon was identified as *M. anatis*. A higher incidence rates were

reported by Chiocco and Bisceglia (1990) who found that the incidence of *M. gallisepticum* from 22 pigeons were 55% and identified them as *M. columbinum* (92.5%), *M. columborale* (65%) and *M. columbinasale* (15%). Moreover, higher prevalence rates of mycoplasmas were mentioned by the findings of Reece et al. (1986B), Astorga et al. (1994), Cookson and Shivaprasad (1994) and Goldberg et al. (1995).

Concerning the use of ELISA for the serodiagnosis of the natural infection of quails with both M. gallisepticum and M. synoviae, the recorded data proved that, for the mean M. gallisepticum titre was 105 and Geometric mean titre (0) under the mean titre group Zero (0-499), while in egg yolk of quails its mean titre was (324) and geometric mean titre (194), under the mean titre of group (0). While the mean M. synoviae titre in sera was (19) and geometric mean titre (2) which layed under the mean titre group "zero", and it was (79) in quail eggs with geometric mean titre (16) which layed under the mean titre of group "zero". These results agreed with those reported by Ansari et al. (1982) who proved that ELISA was more sensitive tool than the (HI) test used to detect antibodies of M. gallisepticum. Morcover, Khokhar and Prasad (1985) who stated that 65 out of 175 serum samples were positive to ELISA, compared with 33 were positive to the (HI) test and 18 were positive to the serum plate agglutination test. Also, Mohammed et al. (1985) proved that no statistical difference between ELISA geometric mean titre of *M. gallisepticum* present in serum and chloroform extracted yolk individual birds. They added that no differences were observed in (HI) or ELISA antibody levels between both egg yolk samples and sera collected samples. Furthermore, Piela et al. (1985) concluded that the chloroform extracted yolks were unsuitable for the use in the HI test for *M. gallisepticum* but produced significant titres with ELISA test. Moreover, Kobisch and Nicolet (1987) reported that the ELISA technique appeared to be the most sensitive procedure recommended to detect early and late antibody titre.

The aformentioned results were contradicted with that reported by Ikoev et al. (1972) who serologically demonstrated specific antibodies to *M. gallisepticum* in Japanese quails proving that such birds were able to contract the natural infection with such microorganisms.

As regard to the results of serodiagnosis of the experimental infection of quails with *M. gallisepticum* by using ELISA technique. The obtained data revealed that the serum samples of quails of the 3rd group, stressed by both environmental and biological stress, gave no detectable specific M. gallisepticum antibodies indicating that such birds did not succumb to the infection. These results agreed with those reported by Yoder and Hofstad (1964) who stated that non of 12 quails inoculated with the first yolk passage of *M. gallisepticum* developed evidence of infection or detectable antibodies. Moreover, Talkington et al.

(1985) mentioned that ELISA test was found to be less sensitive but more specific than S.P.A. less and more sensitive than (HI) test in the detection of antibodies against experimental M. gallisepti. cum.

The considerable gap was present between the obtained findings and that reported by Avakian et al. (1988) who found that the antibodies detected in chickens after experimental infection with M. gal. lisepticum in the first three weeks were better to be detected by S.P.A. test than by ELISA or HI test.

With a meticulous vision one can conclude that the quails, falcons, moorhen fowls and doves as wild birds can act as a reservoir host carrying mycoplasmal organisms. The obtained data revealed that neither such microorganisms nor their specific antibodies could be present, proving that these birds can act as transient carrier for mycoplasmal organisms.

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