Seroprevalence of Coxiella burnetii among humans in Nasiriya

city-South of Iraq.

Jalel abed gati-Msc-Int.Vet.Med.&Preventive – Vet.hospital-thi-qar Abd-ul-aziz salih abd-ul aziz.-Msc.Vet.Micro- college of nursing-Thi-Qar University Alyaa abd-ul-husien Hafeth . M.sc. Immunology. – college of science -Thi-Qar University.

Summary

Out of 146 human(males+females) serum samples, were collected randomly from patients at Nassiriya hospitals from women suffering from abortions and unknown fever which may result from infection with Q fever, they were 46 (31.5) samples seropositive for *Coxiella burnetii*. According to sex there were 13 samples (28.2%) and 33 samples (71.8%) seropositive in male and females respectively. According to age they were 9 (19.59%) seropositive samples in the age 3-20 years, 28 (60.88%) in 21-40 years and 9 (19.59%) samples in 41-89 years The aim of this study is detecting antibodies of phase I and phase II of Coxiella burnetii bacterium, the cause of Q-fever, a zoonotic disease in humans in Thi-qar province (Nasiriyah city-Iraq).

الخلاصة

خضعت (146) عينة مصل جمعت عشوائيا من الإنسان في محافظة ذي قار لفحص الاليزا الذي يكشف خضعت (146) عينة مصل جمعت عشوائيا من الإنسان في محافظة ذي قار لفحص الاليزا الذي يكشف عن أضداد الجرثومة بطوريها ا وπ. بلغت نسبة الإصابة 30.28% ،30.28% في الذكور والإناث على التوالي. أظهرت النتائج وجود (46) عينة موجبة لفحص الاليزا الخاص بالجرثومة وينسبة إصابة بلغت 3.15% في الذكور والإناث ومن جميع الأعمار . واظهرت وجود 13 عينة موجبة (2.82%) من اصل 37 عينة ذكور ووجود 33 عينة موجبة (71.8%) من اصل 109 عينة اناث . كان سبب ارتفاع نسبة الإصابة هو أن العينات البشرية جمعت من أشخاص يعانون من مشاكل صحية مختلفة مشابه لأعراض الإصابة المتوقعة للمرض وذلك يعني ارتفاع مستوى الأضداد في الدم وندرة استخدام المضاد الحيوي تتراسايكلين ومشتقاته ويعد الدواء الاكثر فعالية على الجرثومة. تشير الدراسة الحالية الى انتشار مرض الحمى المجهولة بين البشر في مدينة الناصرية في محافظة ذي قار وتؤكد بان المرض مشتركاً بين الإنسان والحيوان وتوصي بإجراء دراسة موسعة تتضمن تشخيص المرض باستخدام العزل الجرثومي او تفاعل البلمرة وتوصي بإجراء دراسة موسعة تتضمن تشخيص المرض باستخدام العزل الجرثومي او تفاعل البلمرة وتوصي بإجراء دراسة موسعة تتضمن تشخيص المرض باستخدام العرض الحرش واليوان وتوصي بإجراء دراسة موسعة تتضمن تشخيص المرض باستخدام العزل الجرثومي او تفاعل البلمرة والحيوان هدفت الدراسة الحالية إلى الكشف عن أضداد جرثومة المنون العرش واحمى المحبولة في الإنسان والمتوان هدفت الدراسة الحالية إلى الكشف عن أضداد جرثومة المسببة لمرض الحمى المجهولة في الإنسان والحيوان هدفت الدراسة الحالية إلى الكشف عن أضداد جرثومة المسببة لمرض الحمى المجهولة في الإنسان والحيوان هدفت الدراسة الحالية إلى الكشف عن أضداد جرثومة المسببة لمرض الحمى المجهولة في الإنسان

Q.Fever is a zoonotic disease firstontinued risk for infection. The identified in Queensland Australia inotential for transmission is greatly 1935 after an outbreak of febrile illnessnhanced by the extremely low infectious among slaughterhouse workers ⁽¹⁾ theose for *C burnetii*, which is reported to disease was named "Query (Q)" fevere as small as a single organism.⁽¹⁶⁾ because it's etiopathogenesis was not Coxiella burnetii infections have known. In 1935 researchers in the Unitedeen reported in humans, farm animals, States isolated a rickettsial agent fromets, wild animals, and arthropods ⁽⁴⁾. ticks that called nine Mile agent whic Animals are often naturally infected but was subsequently linked to laboratory sually do not show typical symptoms of acquired human infection. $(^{2, 3)}$ the agentSoxiella burnetii infection $^{(3,5)}$. Ticks are were later determined to be identical and onsidered to be the natural primary were eventually named Coxiella burnetteservoir of Coxiella burnetii and in honor of Harold Cox and MacFarlancesponsible for the spread of the infection Burnet. in wild animals and for transmission to

Q fever is a disease caused by adomestic animals ⁽⁴⁾. Cattle, sheep and obligate intracellular bacterium, Coxiellgoats are the main sources of human burnetii. This disease is endemine fection ⁽⁵⁾.

worldwide (3,4) C. burnetii is a gram-Infected animals shed highly stable negative coccobacillus that resides and acteria in urine, feces, milk, and through in host monocytes anplacental and birth fluids. Infection via replicates Macrophages⁽¹⁰⁾ C. burnetii is highlynhalation of aerosolized organisms or infectious; only one organism is requirementation of raw milk or fresh dairy to produce infection under experimental roducts has been reported in humans conditions (11). and animals ⁽⁶⁾. In humans, Q fever is Contact with droplets or fomites manyost often asymptomatic, but acute also result in transmission. Ingestion hasisease (mainly a limited flu-like illness, been proposed as a route of transmission pneumonia or hepatitis) or chronic particularly through the consumption adisease (chronic fatigue syndrome or dairendocarditis) can occur⁽³⁾. Acute Q fever unpasteurized contaminated. products.⁽¹³⁾ however, this mode of a flu-like illness, which is self-limiting transmission is difficult to demonstrator easily treated with antibiotics when an and probably does not play an important propriate diagnosis is made.

role in the spread of disease to Chronic O fever is a severe humans.^(10,14) Sexual transmission of *d*isease that requires prolonged antibiotic has beetherapy because the infection can result I humans burnetii among although this route is endocarditis ⁽⁷⁾ or granulomatous documented, thought to be uncommon.⁽¹⁵⁾ Althoughepatitis ⁽⁸⁾. In addition, the Coxiella direct exposure to parturient animals drurnetii infection can lead to abortions, their birthing products poses the highestillbirth, or pre-mature deliveries in risk for Q fever, the ability of organispregnant women ⁽⁷⁾. In humans, infection to persist in the environment for weeks **results** mainly from inhalation of months after the birth may result in contaminated aerosols from amniotic

fluid, placenta, or contaminated wooMaterials and methods:

but the disease may also be acquired by total of 146 human serum samples (5 the digestive route ⁽⁶⁾. At greatest risk anel of blood inspirited and putted in racks persons in contact with farm animals room temperature to complete clotting (veterinarians, farm workers, butchersThe serum was separated 24 h after as well as laboratory personnel who workampling and stored at -20°C until tested), with infected animals ⁽⁹⁾. In cattle, Qollected randomly from Nasirryia citymainly associated withrag, submitted to ELISA(Enzyme-linked fever is reproductive disorders (abortion, metritismmunosorbent assay) test(Institut and infertility) $^{(17)}$. pourquier), detected antibodies(IgG) of

Routine diagnosis of Q fever ishase I and phase II of Coxiella burnetii usually based on the detection of specificacterium.

by Complement fixation, antibodies

immunofluorescence and enzyme-linkeResults:

immunosorbent assay (ELISA) testOut of 146 human serum samples, Isolation of Coxiella burnetii isollected randomly from patients at hazardous, difficult and time-consumine assiriya hospitals, there were 46 (31.5%) and requires confined biosafety level samples seropositive for Coxiella burnetii laboratories due to the zoonotic nature dfacterium.

the microorganism⁽¹⁸⁾. According to sex there were 13 samples Rapid differentiation of Coxiella burne28.2%) and 33 samples (71.8%)in clinical specimens is very important foropositive females in male and the control of Q fever, because prometpectively (Table 1).

antibiotic therapy may lead to a better

prognosis for individuals (12)

ruble (1) . The percentages of <i>C. but neuri</i> infection in numari.											
sex	male		female	,			Total				
	NO.	%	NO.		%	No.		•	%		
result											
+ve	13	28,2	33	71.8		46		31.5			
-ve	24	24	76	76		100		68.5			
Total	37	25.34	109	74.66		146		100			

Table (1): The percentages of *C. burnetii* infection in human

According to age there were 9 (19.59% According to sex the results showed 35% seropositive samples in the age 3-20 years 2.27% seropositive in males and females 28 (60.88%)in 21-40 years and respectively (Table 2). (19.59%) samples in 41-89 years(Table 2)

Sex	Male		femal	e	Total	
Age	No.	%	NO	%	No.	%
Total samples	37	35.13	109	30.27	146	31.5
(3-20)Years	5	38.46	4	12.12	9	19.59
(21-40)=	7	53.85	21	63.63	28	60.88
(41-89)=	1	7.69	8	24.24	9	19.56
Total +	13	28.2	33	71.8%	46	100%

Table(2) C. burnetii infection related to sex and age in both sex.

Discussion:-

The present study which done in Nasiriya city, as first study, showed that C.burnetii infection is common in our society, out of 146 human samples there are 46 (31.5%)seropositive samples for C.burnetii, this ratio explained the zoonotic and world wide distribution of this microorganism where it is reported in the most area of the world especially in the European , U.k , U.S.A, south America and others.

Our results is similar to many studies in many area such as 48.8% in Ireland farmers(40) and the study which reported in eastern turkey which show that (32.4%) of farmers infection with coxiellosis⁽³³⁾ and the study on Australian abattoir workers which explain that (29%) of workers infection with Q fever⁽³⁴⁾,the study of (234) patients with vascular diseases in bulgaria (34.61%) of them were infection with Q fever⁽³⁵⁾. Basque country (38.5%) , (30%) in ⁽³²⁾, while Sweden farmers different in compared with other ratios such as(13.9%) in $Greece^{(37)}$, (16%) in Australian cattle workers(38),(15.78%) of Bulgarian patients with pneumonia ⁽³⁵, (12.7%) (10.8%) (18.7%) in Madrid, Sevilla and lanzarote respectively(29-24).

This increasing in the prevalence of *C. burnetii*

infections in Iraq is expected because of cumulative exposure and multiple of the reservoirs in the nature which include many species of mammals, birds and ticks also the infections of *C.burnetii* is most often latent in animals with persistent shedding of bacteria to the environment however in females intermittent high level shedding occur at the time of parturition with millions of bacteria being released per gram of placenta fluids ⁽¹⁹⁾.

The other cause of prevalence of *C.burnetii*, is a limit using of tetracycline family which is the drug of choice in treatment of *C.burnetii* infection in human patients in Iraq generally.

Human are usually infected by contaminated aerosols from domestic animals particularly with after contact parturient females and their birth products ⁽²¹⁾. Also the high risk to infection exposure to is increasing by direct contact with animals such as butchers.

veterinarians, framers ⁽²²⁾. Also, the consumption of contaminated foods e.g.: un pasteurized milk and diary products . These risks is common in our society because the nature of lifestyle and direct contact with animals.

from Data seroepidemiological studies conducted in other European countries have shown low prevalence of C.butntii infection 15 % has been found in Italy, Czechoslokia republic of ,Switzerland and Sweden ⁽²⁵⁻²⁶⁾, 19% , 23% in Germany ²⁷⁾ and France respectively ²⁸⁾. These differences in the seroprevalence of C.burnetii in the regions which mention above may due to economy based on fishing and service sector activities (21-20.23).

From the main reasons of differences of previous studies in prevalence of *C.burntii* because this studies based mainly on serologic tests especially (ELISA), Complement fixation and indirect immunoflourescent assay however the seroprevalence of *C.burnetii* infections various widely from <u>1</u> country to another and from <u>1</u> state to another and also the reports from same state show wide differences depending on testing methods and year of surveys $^{(29)}$

The results of this study shows that the percentage of infections of Q.fever in men (35.13%) is more than the percentage in women (30.27%) (Table 3) and this result explain that the prevalence of *C.burnetii* infections was higher in males than woman as reported in other studies and this probably because Q. fever is an occupational diseases⁽³⁰⁾.

According to the (Table 2) which include the percentage ratio of *C.burnetii* infection according to the age (in both males and females) note that the high infection is concentrate in adult ages (21-40) as (53.85%) , (63.63%) in males and females respectively and this ratios is corresponding with most studies which showed that the high ratios of infection with Q fever occur in the adult ages^{.(31)}. The ages (4-67) years old with the median of 43.5 old are mostly associated with Q fever infections.⁽³⁸⁾.

In conclusion, the result of this study confirm the prevalence of anti- C.burnetii antibodies in Nasiriya region, the dada obtain from this study may be useful as reference in further studies in thiqar region or in Iraq. Further studies in Iraq, beside the collaboration between veterinary and medical services on coxiella infection in both domestic animals and human are needed to elucidate the epidemiology of Q fever in Iraq.

References :-

1-Derrick, E.H. (1937). "Q" fever, new fever entity: clinical features, diagnosis, and laboratory investigation. Med J Aust.;2:Pp281-299. 2-Davis, G.E.and Cox, H.R(1938).A filter-passing

infectious agent isolated from ticks. I. Isolation from Dermacentor and ersoni, reactins animals. and filtration in experiments. Public Health Rep;53:Pp2259-2261. 3-Maurin, M., Raoult, D. (1999). Q fever. Clin. Microbiol. Rev.,; 12: Pp518-553. 4- Norlander, L. (2000). Q fever epidemiology and pathogenesis. Microbes Infect.,; 2:Pp 417-424. 5-Hilbink, F., Penrose, M., Kovacova, E., Kazar, J. (1993). Q absent from New fever is Zealand. Int. J. Epidemiol.,; 22: Pp945-949. 6- Fishbein, D.B. and Raoult, D. (1992). A cluster of Coxiella burnetii infections associated with exposure to vaccinated goats and their dairy products. unpasteurized Am. J. Trop. Med. Hyg.,; 47:Pp35-40. 7- Raoult, D. and Marrie, T.S. (1995). Q. Fever., Clin. Infect. Dis.,; 20:Pp 489-495. 8- Weir, W.R.C., Bannister, B., Chambres, S., De Cock, K., Mistry. H. (1984). Chronic Q fever associated with granulomatous hepatitis. J. Infect.; 8: Pp 56-60 9- Hatchette, T.F., Hudson, R.C., Schlech, W.F., Campbell, N.A., Hatchette, J.E., Ratnam, S., D., Donovan, C., Raoult. (2001). Marrie, T.J. Goatassociated Q fever: a new disease in Newfoundland.Emerg. Infect. Dis.,; 7: Pp 413-419.

10. Maurin, M. and Raoult, D.(1999). Q fever. Clin Microbiol *Rev*;12: Pp 518–553.

11. McCaul, T.F. and Williams, J.C.(1981). Developmental cycle of *Coxiella burnetii*: structure and morphogenesis of vegetative and sporogenic differentiations. J Bacteriol;147:1063–1076.

12- Zhang, G.Q., Hotta, A., Mizutani, M., Ho, T., Yamaguchi, T., Fukushi, H., Hirai, K. (1998). Direct identification of Coxiella burnetii

plasmids in human sera by nested PCR. J. Clin. Microbiol.;36: Pp 2210-2213.

13. Lennette E.H, Clark W.H, Abinanti M.M, et al. (1952). The effect of pasteurization on *Coxiella burnetii* in naturally infected milk. Am J .Hyg;55:Pp 246–253.

14. Krumbiegel, E.R. and Wisniewski, H.J. (1970).Q fever in Milwaukee. II.Consumption of infected raw milk by human volunteers. Arch Environ Health;21: Pp63–65.

15. Milazzo, A. Hall R, Storm PA, et al. (2001). Sexually transmitted Q

fever. Clin Infect Dis;33:399–402.

16. Tigertt, W.D, Benenson, A.S, Gochenour, W.S. (1961). Airborne Q fever. Bacteriol Rev;25: Pp 285–293.

17- Hissig, M., and Lubsen, J. (1998). Relationship between abortions and seroprevalences to selected infectious agents in dairy cows. J. Vet.

Med. B.,; 45: Pp 435-441.

18- Scott, G.H. and Williams, J.C. (1990). Susceptibility of *Coxiella burnetii* to chemical disinfectants. Ann N Y Acad Sci;590:Pp 291–296.

19- Tringali, G. and Mansueto, S.(1987). Epidemiology of Q fever in Italy and other miditerrranean counteries. Zbl Baki Hyg A;267:20-25.

20- Sanzo JM, Garda-Calabuig MA, Audicana A, Dehesa V. Q.(1993). fever prevalence of antibodies to *Coxiella bumetii* in the Basque country, Int J Epidemiol.:22:Pp 1183-88.

21- maurin, M. and Raoult,D. (1999). Q.FEVER.clin microbial Rev, oct;12(4): Pp 518-53.

22- TraUero E, Cilia G, Montes M, Saenz-Dominguez JR, Alcorta M. (1995).Prevalence of *Coxiella bumetii* infection among slaughterhouse workers in northern Spain. Eur 3 Clin Microbiol Infect Dis;14: Pp71-73.

23-Tellez, A. and Martin, A.(1999). Study of coxiella burnetii human and animal seroprevalence in rural population in madrid community.Eur J Epidemiol :5:Pp 444-46.

24- Pascual-Velasco F, Rodriguez-Perez JC., Otero, I., Borobio MV. (1992). Seroprevalenaa de la fiebre Q en la poblad6n adulta de Lanzarote(Islas Canarias). An Med Interna:9: Pp 428-32.

25- Tringali, G. and Mansueto, S. (1987). Epidemiology of Q fever in Italy and other Mediterranean countries. Zbl Bakt Hyg *A*:267:Pp20-25.

26- Macellaro, A., Akesson, A., Norlander, L. (1993). A survey of Q-fever in Sweden. Eur J Epidemiol;9: Pp 213-16.

27- Heinrich, R., Naujoks-Heinrich, S., Saebisch, R. (1983). Seroprevalence of Q fever in an endemic area of Southern Germany. Dtsch Med Wochenschr,108:Pp 1318-24.

28- Raoult, D., Drancoun, M., De Micro, C. (1986).Les hepatitis de la fievre Q. A propos de 14 cas et revue de la literature. Sem Hop Pans:62: 997.

29- wisniewski, H. and krumbigel, E. (1970). Q Fever in the Milwaukee area. Arch Enveron Health.;21: Pp 58-62.

30- pascual –velasco, F., Montes, M., Marimon, JM., Gilla. C. (1998).High seroprevalevce of coxiella burnetii infection Eastren in cantabria(spain). International journal epidemiology.;27:Pp 142-145.

31- Raot, D. and Marrie, T. (1995). Q.fever. clini infect Dis.;20: Pp 489-96.

32-A. Macellaro, A. Akesson² and L. Norlander(1993).A survey of Q-fever in Sweden fever in Sweden Journal European Journal of Epidemiology Issue Vol. 9, No. 2 / March. 33-

Senay,S;Zulal,O;Ufuk,D;Biray,O .(2006). The seroprevalevce of coxiellosis in farmers and cattle in Erzurm District in Turkey.Institution of vetrenary control and researches.Erzurm.Turkey.Jurk.J. Vet.Anim.Sci.30: Pp 71-75

34- Gilroy N, Formica N, Beers M, et al. (2001). "Abattoirassociated Q fever: a Q fever outbreak during a Q fever vaccination program." *Australian and New Zealand Journal of Public Health* 25(4): 362-367.

35- Martinov.S.(2007). Contemporary state of the problem Q fever in Bulgaria.National Research Veterinary Medical Institute,Sofia, Bulgaria.

36- Papadogiannakis1 E.*, Kontos1 V., Kontou1, I. Kostomitsopoulos2, N. Siochou1, E.. Tsachev3,I Vassalou1, E. Makropoulos, V. (2007) . A SEROLOGICAL SURVEY OF BRUCELLOSIS,ECHINOCOCC OSIS,QFEVER,TOXOPLASMO SIS,

LEISHMANIASISANDMEDIT ERRANEANSPOTTEDFEVERI NANIMAL

PRODUCTIONEMPLOYEES

IN GREECE. *Trakia Journal of Sciences*, Vol. 5, No. 2, Pp 70-78,

37- Shapiro R, Siskind V, Schofield F, et al. (1990). "A randomized, controlled,doubleblind, cross-over, clinical trial of Q fever vaccine in

selectedQueensland abattoirs." Epidemiology Infection and 104(2): Pp 267-273. 38-Costa.p: Brigatte,M; Greco, D. (2006). Quasting one brazillian Query: Reporting 16 cases of Q .fever from Gerias, Brazil. Med. J.50:Pp 333-338. 39- McCaughey. C, McKenna .J , McKenna. C , P. V. Coyle, H. J. O'Neill , D. E. Wyatt , B. Smyth and L. J. Murray (2008).

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