SERO - EPIDEMIOLOGY OF SHEEP BRUCELLOSIS IN BAGHDAD.

*K.A. AL - Salih *J.M. Kalaf **A.H. Jawad ***G.M. Ebrahim

*Department of Veterinary Medicine and Therapeutics, College of Veterinary Medicine, University of Baghdad
**Communicable Disease Center (CDC), Baghdad, Iraq.
***Public Health Center, Baghdad, Iraq.
(Received 15 April 2005, Accepted 23 January 2006)

Keywords: Serum, IFAT, Brucellosis

ABSTRACT

A total of 531 serum samples were subjected to Rose Bengal test (RBT), serum agglutination test (SAT) and indirect fluorescent antibody test (IFAT) and ELISA. The incidence of brucellosis was found to be 55.76%, 47.64%, 55.76% and 58.70% on basis of RBT, SAT, IFAT and ELISA respectively. The agreement between RBT and SAT was 85.47% and between RBT and IFAT was 100% giving a positive result. In case of ELISA the positive results were higher and we recommended using it as the definitive test for detection of brucellosis.

INTRODUCTION

Brucellosis is one of the zoonotic diseases and is a problem of both public health and economic importance in several countries of the world, and is caused by bacteria of the genus Brucella which infect all domestic animals as well as human beings. It causes abortion and infertility in animals (10, 12) and undulant fever in man (3). In Iraq it is an enzootic disease (2) and there are few studies on the incidence of brucellosis in sheep have been carried out (20, 17, 16).

There are many serological tests used for diagnosis like RBT, SAT (5, 6) but they may give false negative or false positive (9) especially SAT which cannot differentiate classes of immunoglobulins (7). It is not always possible to isolate the causal organism from infected animals, so the serological tests play a major role in the routine diagnosis of brucellosis (5). Now anew technique was used to prevent many problems occur during many serological tests. This technique is ELISA which have the higher sensitivity to diagnosis of brucellosis (6, 2, 8).

The present study was conducted to find the incidence of brucellosis in sheep in three animal flocks in Baghdad by using four serological tests, Rose Bengal test (RBT), serum tube agglutination tests (SAT), Indirect fluorescent antibody test (IFAT) and Enzyme-Linked Immunosorbent assay (ELISA).

MATERIAL AND METHODS

Studies were performed on (531) blood samples collected from sheep in three separated flocks with history of abortion in Baghdad. Serum samples were tested by the previous serological antibodies by IFAT according to (8). All positive serum samples was tested by SAT as described by (5). IFAT also was carried out as described by (1), and ELISA by using antigen produced by Diasorin company and IgA indicator that contain Brucella spp. The method was done according to Diasorin company instructions.

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RESULTS

1- Rose bengal test (RBT):- A total of 913 blood samples were collected from three flocks examined by this test to detect the presence of brucella antibodies. The results revealed 296 (55.76%) positive reactors out of total number of the test samples as shown in table 1. While the number and percentage of positive reactors serum in the test flocks 120 (58.53%) 80 (51.47%) and 96 (56.47%). In first, second and third flocks respectively. The remaining 235 (44.34%) were negative to RBT.

2- Serum tube agglutination test (SAT):- All positive reactors serum samples in RBT were examined by SAT. The results as shown in table 2 revealed 75, 37, 45, 63, 33 positive reactors in titer 1/10, 1/20, 1/40, 1/80, 1/160 respectively while 43 positive reactors sample in RBT were negative in SAT. The agreement between the RBT and SAT was (85.47%) in giving positive results.

3- Indirect Immunofluorescent antibody test: - The same positive reactors serum samples in RBT were examined by IFAT. The result as shown in table 2 revealed 28, 51, 50, 116 positively reactor in titer 1/16, 1/32, 1/64, 1/128 and 1/256 respectively. The agreement between RBT and IFAT was (100%) in giving a positive result.

4- All positive and negative samples in RBT were examined by ELISA. The results show that 29 samples from those which were negative in RBT give positive reaction and 15 sample of those with positive RBT were negative in ELISA. The reading 0.208 was the limit between positive result in ELISA. The agreement between IFAT and ELISA was 100%.

DISCUSSION

The incidence of brucellosis in sheep in the examined flocks was 55.76% as screened by the RBT. This result varies significantly from those reported by (20, 17, 16, 3) who found that the incidence of brucellosis in sheep were 0.1%, 1.1%, 0.93% and 7.91% respectively. The high incidence of brucellosis (55.76%) in this study was also reported by previous studies in different provinces of Iraq (12). While (14) have noticed an incidence averaged out to 20% in aborted and non-abortion goats in different provinces of Iraq. Of 296 sera positive to RBT 253 were positive to the SAT giving an agreement of (85.47%) between the two tests. This result was comparable with result reported previously by several workers (16, 3). (18) whom inoculated six sheep with B. melitensis strain H 38 and showed that RBT identified infected animals in earlier stage than the SAT and was often positive when the SAT was negative or inconclusive. There was marked drop in the agglutination test titer in 34 weeks after challenge. While the RBT was still showing positive reaction. Concerning IFAT results obtained during this study it was shown that all positive reactors sera in RBT were positive in IFAT, giving an agreement 100% between the two tests and high number of serum samples (116) giving high titer 1/256.

The difference in serological response to the three tests may be due to the fact that the antibody classes mediating these tests may differ in sheep and the length of time that animal remain infected as a reservoir host, may involve different classes of antibody reacting in various tests (22). It must inevitable include some animals with negative serology and other with titers at every up to the maximum possible, this is due to the fact that after infection, the antibody titers in various serological tests will vary with the course of antibody production (5).
During the incubation period, the results of serological tests may be negative although infected animals may abort soon after wards.

The result of the SAT is either negative or inconclusive in the incubation stage or in the late chronic stage of brucellosis, while RBT seems to identify infected animals at an earlier stage than the SAT and is often positive when the SAT is negative or inconclusive but IFAT is positive (21,18).

This may be ascribed to the type and concentration of immunoglobulin induced in animals due to brucella infection, so IFAT was proven to detect both an initially induced antibodies early infection and late induced antibody (late stage of infection) in brucella infection. These results are compared with those reported by (3), they used IFAT in diagnosis of human brucellosis and they concluded that IFAT are very reliable and sensitive test for diagnosis of brucellosis.

With regard to the incomplete agreement of ELISA and RBT test in this study we observed that this may occur due to the positive reaction with non-specific antibodies which give false positive reaction (4). The false negative results may occur due to the low degree of sensitivity against immunoglobulin because the high sensitivity is against IgM with comparison to IgG(4,11).

In case of SAT and its little agreement with ELISA may be due to the early and late stages of infection that give false negative in SAT (19,15). In this study we can conclude that RBT is a good serological test in case of surveys but ELISA and IFAT were the most accurate which make them the most dependable tests in diagnosis and special research on brucellosis. SAT must be under more studies to evaluate its capability to used as a diagnosis tool in brucellosis.

Table 2- Show titer of positive serum with RBT-in tube agglutination test and Immuno- Fluorescent antibody technique.

<table>
<thead>
<tr>
<th>No of samples</th>
<th>Titer in tube agglutination</th>
<th>Titer in Immuno Fluorescent antibody technique</th>
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<tbody>
<tr>
<td></td>
<td>10</td>
<td>20</td>
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<tr>
<td>120</td>
<td>84</td>
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<td>80</td>
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<td>19</td>
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</table>
دراسة وزيادة ومصلية عن البروسيلا في الأغنام في بغداد

كريم عاكلس، مسئول مجمع عادلة، خبراء في مجال الألبستر، بناءة، وبحث وتفتيح، الطبي، البصري-كلية الطب البيطري، بغداد، العراق

مركز الإستراح المتواضعة، بغداد، العراق

مختبر الصحة المركزية، بغداد، العراق

الخلاصة

(ROSE Bengal test) اكتشاف القيمة 296 من النتائج. الدراسة، التعبير عن تفاعل الألبستير ( ở انحلال) لمصل (tube agglutination test)، والتفاعل (ELIZA) (fluorescent antibody test).

عند اجراء اختبارات البروسيلا، تكاثر، في المصل الإلبيوي، اختصار الاستماع البيولوجي، الفيروسات والكواترا على التوالي، وكان التوافق بين اختبار البروسيلا، تكاثر المصل الإلبيوي، في 87%، 85%، 83%، 81%، 79% من اختبارات البروسيلا، تكاثر المصل الإلبيوي.

في 99% من اختبارات الفيروسات، مصل، البروسيلا، في 99% من اختبارات الفيروسات، مصل، البروسيلا.

للمزيد، تم استخدام معالجات مبكرة للعظام لتفعيل نشاط أنتي-gx, ump,q,

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