Serologic Survey of Crimean-Congo Haemorrhagic Fever among Sheep in Ardabil Province, Northwest Iran

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INTRODUCTION

Crimean-Congo Hemorrhagic Fever (CCHF) is a vector-borne viral zoonotic disease, which causes acute febrile illness in humans. The disease in humans is often accompanied by hemorrhage, with the fatality rate ranging 5 to 30%. The causing virus is usually transmitted to humans by tick bites or direct contact with the blood or tissues of a viremic animal [1, 2]. *Hyalomma* spp. ticks are considered to be the most important vector of the virus, determining the distribution of CCHF virus worldwide [3]. A wide variety of livestock (sheep, cattle, and goats) and wild animals can be infected with this virus, but sheep are considered to be the most important domestic host for the virus in nature [1]. The infection is usually asymptomatic and subclinical in animals [2]. Slaughterhouse workers, butchers, farmers, veterinarians, and shepherds are at higher risk of infection [4]. Human cases of CCHF have been reported from different parts of Asia, Africa, Europe and the Middle East [2].

In Iran, the existence of CCHF among animals was first reported in 1970 [5]. In 1999, the first cases of human infection with CCHF were reported in Shahrekord City (central Iran) and subsequently in other provinces of the country [6]. Human infections of CCHF has been reported from most of the provinces [7], and Human cases of CCHF are very few in some western and northwestern provinces of Iran [8, 9].

There is not much data on the status and distribution of CCHF infection among domestic animals in Ardabil Province. The data on livestock in this area is limited to a single study of animals (cows, sheep, and goats) in Meshkin-Shah County (central Ardabil) in 2002-2005, in which the seroprevalence of CCHF was 39.3% [7]. The present study was conducted to investigate CCHF status among sheep in different geographical regions of Ardabil Province.

Keywords: CCHF, Seropositivity, Epidemiology, Zoonosis, Iran.

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MATERIAL AND METHODS

Study area and sampling. This study was carried out in 2011. Ardabil Province is located in northwestern Iran with an area of approximately 17953 Km². It borders the Republic of Azerbaijan in north and northeast, East Azerbaijan in the west, Zanjan in the south, and Gilan in east. Based on a report by the Iranian veterinary organization (2009, unpublished data), this province has 2,316,990 sheep, 262,190 goats and 348,150 head of cattle.

In this study, the blood samples of sheep were collected randomly from the regions in the north (Parsabad and Bilasavar counties), center (Ardabil, Nir, Meshkin-Shahr counties) and south (Khalkhal and Kowsar counties) of the province. We included 29 herds in the study and recorded their specifications including age, gender, and the habitats. Samples were sent immediately to the laboratory, and the sera were separated and stored in -20°C until used.

Serology. Serum samples were analyzed by a specific ELISA for detection of IgG antibodies. The process involved coating the ELISA plates with mouse hyperimmune ascitic fluid (diluted at 1:1000) in phosphate-buffered saline (PBS 1x) and incubation overnight at 4°C. Following the washing step, the native or recombinant antigen in PBS (diluted at 1:500) containing 0.5% Tween (PBST) and 3% skimmed milk (PBSTM) was added, and the plates were incubated for 3 h at 37°C.

The serum dilutions in PBSTM (1:100) were added, and the plates were incubated for 1 h at 37°C. Peroxide-labelled anti-animal immunoglobulin diluted at 1:350 in PBSTM was added, and the plates were incubated for 1 h at 37°C. The plates were washed 3 times with PBST after each incubation. Finally, hydrogen peroxide (H2O2) and 3, 3’ , 5, 5’ tetramethylbenzidine (TMB) were added, and the plates were incubated for 15 min at room temperature. The enzymatic reaction was stopped by the addition of 4 N H2SO4. The plates were read by the ELISA reader (Anathos 2020) at 450 nm [10, 11].

Data Analysis. The data were analyzed using the SPSS for Windows statistical package (SPSS Inc., Chicago, IL, 16th version). Chi-square test was used for the comparison of variables in the analysis and the P-values less than 0.05 were considered statistically significant. ArcGIS software (ESRI, version 9.3) was used for mapping the results.

RESULTS

In this study, 256 sheep sera belonging to 36 (14.1%) rams, and 220 (83.9%) ewes were obtained from the northern (66), central (120) and southern (70) regions of Ardabil Province. The average (=SE) age of the animals was 4.38±0.16 years.

The seroprevalence of anti-CCHF IgG antibodies in sheep was 27.34% (95% CI: 22.25% - 33.11%) (Table 1). Although the seropositivity rates of CCHF in the northern regions (36.36%) was higher than in the central (26.27%) and southern (20%) regions, the difference was not statistically significant (p=0.10). The highest seropositivity for CCHF was detected in Parsabad County (45.57%), and the lowest was in Khalkhal (17.78 %) (Figure 1).

No significant difference in seroprevalence rate of the CCHF was observed between rams (36.11%) and ewes (25.90%) (p=0.20). Neither, no significant difference was observed between different age groups and the seroprevalence rate of CCHF (p=0.54).

DISCUSSION

In this study, 27.35% of sheep from various regions of Ardabil Province showed a history of CCHF infection. In Iran, CCHF seroprevalence rate in sheep showed a broad range in different areas, 77.5% in three Khorasan Provinces (northeastern Iran, 2003-2005) [12], 3.7% in Mazandaran Province (northern Iran, 2010-2011) [13], 76.9% in Isfahan Province (central Iran, 2002) [14], 27.4% in Bahar County of Hamadan Province (western Iran, 2006) [15], and 15.6% in Eastern Azerbaijan Province (northwestern Iran, 2010) [16]. In a study conducted between 2004 and 2005 in Meshkin-Shahr County of Ardabil Province, 39.3% of livestock (41.9% sheep, 33.3% goats, 30.0% cattle) had a history of CCHF infection [7], this rate of seropositivity was higher than what we observed in this study (25%).

It has been shown that CCHF antibodies (IgG) can persist for more than 5 years in animals blood [17] and various studies have observed a positive association between the age of the animals and the disease seroprevalence [13, 16, 18]. In this study, however, no association between age and the rate of seropositivity was noted. Furthermore, in agreement with previous studies carried out in Eastern Azerbaijan [16] and Mazandaran [13], no significant relationship between gender and seropositivity was noticed in this study.

The relatively high CCHF seroprevalence among sheep in this study will strengthen the possibility that further cases of human disease might have occurred in this province without being diagnosed. One other probable hypothesis is that the CCHF virus strain circulating in these regions does not have a high pathogenicity for humans.

Table 1. Seropositivity rate of CCHF among sheep in different geographical regions of Ardabil Province

<table>
<thead>
<tr>
<th>Region</th>
<th>Number tested (% Seropositive)</th>
<th>County name</th>
<th>Number tested (% Seropositive)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Northern</td>
<td>66 (36.36)</td>
<td>Parsabad</td>
<td>35 (48.57)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Bilasavar</td>
<td>31 (22.58)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ardabil</td>
<td>54 (29.63)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nir</td>
<td>30 (23.33)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Meshkin-Shahr</td>
<td>36 (25.00)</td>
</tr>
<tr>
<td>Central</td>
<td>120 (26.67)</td>
<td>Khalkhal</td>
<td>45 (17.78)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Kowsar</td>
<td>25 (24.00)</td>
</tr>
<tr>
<td>Southern</td>
<td>70 (20.00)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>256 (27.34)</td>
<td></td>
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</tbody>
</table>
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CONFLICT OF INTEREST

The authors declare that there are no conflicts of interest associated with this manuscript.

REFERENCES


