Detection of *Neospora caninum* IgG Antibodies in Goats in Elazig, Erzurum and Kırsehir Provinces of Turkey

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**ABSTRACT**

The aim of this study was to detect *Neospora caninum* IgG antibodies in Saanen and Hair goats from Elazig, Erzurum and Kırsehir provinces of Turkey. For this, a total of 128 sera obtained from goats were tested using a commercially available competitive enzyme-linked immunosorbent assay (c-ELISA) kit. Overall prevalence of anti-*N. caninum* antibodies was 10.2% (13/128); and mean prevalence in Hair and Saanen goats were 13.8% (12/87) and 2.4% (1/41) respectively. This is the first serological study on caprine neosporosis in the above mentioned provinces of Turkey.

**Keywords:** *Neospora caninum*, goat, c-ELISA, Turkey.

**INTRODUCTION**

*Neospora caninum* is a coccidian parasite. It was first identified in dogs with encephalomyelitis and myositis in 1984, and was described as a new genus and species in 1988. Dogs are both definitive and intermediate hosts. Cattle, sheep, goats, horses and deer are intermediate hosts. Dogs can acquire infection by ingestion of infected tissues; and intermediate hosts can be infected either by horizontal postnatal infection or by vertical transmission during pregnancy (1-3).

In *Neospora* infection, cows can abort; fetuses may die *in utero*, be resorbed, mummified, autolyzed, stillborn, born alive with clinical signs or born normally, but persistently infected. Moreover, infection may cause premature culling, diminished milk production and repeat breeder problems in herds (1-7). Although neosporosis is a major problem in cattle, serological, molecular and experimental data show that *N. caninum* can cause clinical infections in goats. Abortions, fetal deaths and stillbirths have been reported in goats due to *N. caninum* (8-13). *N. caninum* antibodies were reported in goats in Southern and Northern Jordan, Poland, Brazil, Sri Lanka, Taiwan and Turkey (14-21). The aim of this study was to detect *N. caninum* IgG antibodies in Saanen and Hair goats from Elazig, Erzurum and Kırsehir provinces of Turkey.

**MATERIALS AND METHODS**

**Sample collection**

Blood samples were collected from the jugular vein in sterile tubes from 128 dairy goats originating from Saanen and Hair breeds in Erzurum, Elazig and Kırsehir provinces of Turkey (Figure 1). Age of the animals ranged from 2 to 4 years. Sera were removed after centrifugation at 2000 rpm for 5 minutes and stored at -20 °C until tested.

**Serologic examination**

Antibodies to *N. caninum* were detected using a commercially available competitive enzyme-linked immunosorbent assay (c-ELISA) kit (VMRD, USA). The test was carried...
out by following the instructions of the manufacturer. The mean optical density (OD) at 630 nm was determined for all wells using a microplate reader (ELx 800 UV, Universal Microplate Reader, Bio-Tec Instruments, Inc., VT, USA). The percent inhibition for each test sample was determined using the formula:

\[
\text{Inhibition (\%)} = 100 - \left( \frac{\text{Sample O.D.} \times 100}{\text{Mean Negative Control O.D.}} \right)
\]

The samples with values of ≥ 30% inhibition were regarded as positive and those with the values < 30% inhibition were regarded as negative.

**Data analysis**

Data management and statistical analysis by Fisher’s exact chi square tests were performed using SPSS 10.1 software for Windows. The level of significance was considered to be 5% (P<0.05).

**RESULTS**

The overall seroprevalence of *N. caninum* was 10.2% (13/128). Mean seroprevalences of Hair and Saanen breeds were 13.8% (12/87) and 2.4% (1/41) respectively. No statistical difference were found between Saanen and Hair breeds (P>0.05). Results of the study are summarized in table (Table 1).

**DISCUSSION**

Neosporosis is a major problem for cattle breeders, and causes economic loses in the dairy industry throughout the world (1-3). Therefore, existing studies are concentrated on bovine neosporosis; and there is limited serological data about caprine neosporosis (1-3). Abo-Shehada and Abu-Halaweh studied 302 goats from 62 flocks in northern Jordan, and found the prevalence to be 12% at flock level and 2% at the individual level, respectively (14), Al-Majali *et al.* reported individual level and flock level seroprevalence as 5.7% and 48.7% in 300 goats from 24 flocks in southern Jordan (15), Czopowicz *et al.* tested 1060 sera for antibodies against 256	256

Figure 1: Location of study areas in Turkey. Areas shaded in red are covered in this study. Grey-shaded areas have been explored in previous studies.

![Map of Turkey showing study areas](https://example.com/map.png)

Table 1: Seroprevalence of *N. caninum* in different provinces of Turkey.

<table>
<thead>
<tr>
<th>Location</th>
<th>Examined (n)</th>
<th>Infected (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Saanen</td>
<td>Hair</td>
</tr>
<tr>
<td>Elazig</td>
<td>- 70</td>
<td>- 70</td>
</tr>
<tr>
<td>Erzurum</td>
<td>- 17</td>
<td>- 17</td>
</tr>
<tr>
<td>Kırsehir</td>
<td>41</td>
<td>- 41</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td>41</td>
<td>87</td>
</tr>
</tbody>
</table>
Neospora caninum and determined the true herd level prevalence as 9.0% in Poland (16). Uzdža et al. found the prevalence as 15% in 385 goats in Bahia, Brazil (18), Faria et al. found the prevalence as 3.3% in 306 goats in the north-east region of Brazil (17), other researchers reported the seroprevalence rates as 7.0% (3/486) in Sri Lanka (19) and 0% (0/24) in Taiwan (22). Different diagnostic tests such as ELISA, indirect fluorescence antibody test (IFAT) and enzyme immuno-assay (EIA) were used in the reported studies (14-19, 22).

Seroprevalences of Neospora caninum were determined to be 25.9% (47/181) in Nigde, and 5% (9/180) in the Sanliurfa provinces of Turkey (Figure 1) (20, 21). In this study, we identified the overall prevalence as 10.2% (13/128) in three different provinces of Turkey. Mean seroprevalences were 23.5%, 11.4%, and 2.4% in Erzurum, Elazig and Kırşehir provinces, respectively. Seroprevalence rates decreased from east to west. In addition, mean seroprevalence of Hair and Saanen goats were 13.8 % and 2.4 %. Although the test (c-ELISA) used and the sex of animals tested was the same, seroprevalence rates were different in three separate studies conducted in five different provinces of Turkey (20, 21). In these studies, there is no data about the breeding and feeding systems of goats. In their study, Cayvaz and Karatepe did not mention the breed of goats, and ages of the animals ranged from 1 to 5 years (20). Sevgili et al. studied Aleppo and Hair goats between the ages of 1-6 years (21). In the present study, we detected Neospora caninum antibodies in Saanen and Hair goats between the ages of 2-4 years. Study areas of three research projects were Central Anatolia (Nigde and Kırşehir), East Anatolia (Elazig and Erzurum) and Southeastern Anatolia (Urfa) which have steppe, continental and semi-arid continental climate, respectively. Southern Anatolia is the driest and East Anatolia is the coldest regions of Turkey (23). Likewise, seroprevalence rates are different in two different parts of Jordan and Brazil (14, 15, 17, 18). Therefore, these differences can be explained by the use of different breeds, sample sizes, climatic factors and breeding and feeding systems.

At present, there is no effective treatment for neosporosis (1). For prevention against caprine neosporosis, routine serological surveys can be done in goat flocks; and in case of low seroprevalence rates, positive dams can be culled. Reduction in the population of stray dogs and limiting their access to goat food, water sources, or to placental membranes and carcasses of aborted goat fetuses may control postnatal infection and decrease the risk of contamination of the environment and later infection with oocysts shed in the feces of dogs.

In the present study we determined the rate of exposure of caprine neosporosis in Erzurum, Elazig and Kırşehir provinces of Turkey. We consider that further serological and clinical studies on caprine neosporosis are necessary for developing effective control programs. A clear understanding of the epidemiology of the disease both in Turkey and globally is required.

REFERENCES


