

STUDIES ON THE PREVALENCE OF ANTIBODIES FOR PESTE DES PETITS RUMINANTS (PPR) AND RINDERPEST (RP) VIRUS IN SHEEP AND GOATS SERA

BY

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SUMMARY

A total of 182 sera samples were collected from two sheep breeds (mixed and Barki sheep) as well as goats raised under arid conditions and examined by serum neutralization test (SNT) for neutralizing antibodies of PPR and RP viruses.

The results showed that at 1:4 serum dilution, 29.41% of mixed breed sheep, 29.82% of Barki sheep and 29.82% of goats sera contained antibodies to PPR virus, while 2.9%, 12.28% of the two sheep breeds respectively and 7.0% of goats sera have antibodies to RP virus.

It could be concluded that the prevalence of neutralizing antibodies to PPR virus is equal in sheep and goats. No significant difference between the two sheep breeds tested for PPR virus antibodies, but RP virus antibodies was higher in the sera of Barki sheep than that of the mixed breed sheep. and the antibodies for PPR virus in the two sheep breeds were more prevalent than in goats.

INTRODUCTION

Peste des petits Ruminants (PPR) is a virus disease of goats and sheep that characterized by fever, nasal and oral discharge and mucosal erosions, often followed by severe enteritis and pneumonia. The disease is distinct from that caused by rinderpest virus in goats and sheep (Lefevre, 1982). Pulmonary involvement predominate in PPR whereas digestive lesions are dominants in rinderpest.

PPR virus has been considered as a variant of rinderpest virus naturally adapted to cause disease in goats and sheep but not cattle (Mornet et al, 1956). Subsequent studies of Gibbs et al., (1979) confirmed the antigenic difference between PPR and RP virus. The two viruses also are closely related serologically but are not identical and cross neutralization differentiate between them (Hamdy et al., 1976). PPR virus protected cattle against rinderpest and RP virus protected sheep and goats against PPR virus (Gibbs et al. (1979) and PPR virus was proposed as a fourth member of the Morbilli virus group.

In Egypt PPR virus was isolated from goats by Ismail and House (1990) and from sheep by Fayed et al., (1990). The pathogenicity of the virus for goats was studied by Ismail et al., (1990) and neutralizing antibodies in a percentage of 28.97 of sheep sera was reported by Hassan (1994).

The aim of this work was to study the prevalence of neutralizing antibodies in sheep breeds as well as goats for both PPR and RP virus.

MATERIAL AND METHODS

Blood samples:

A total of 182 blood samples were collected from 68 mixed breed (Barki x Marino), and 57 Barki sheep as well as 67 goats raised at the Desert Research institute station at Mariott. The sera were separated and kept at -20 C until examined.

Viruses:

PPR virus strain "Egypt 87" was obtained from the virus strains collection bank of Animal Health Research Institute, and RP virus live attenuated "Kabete O" strain from Abbassia Serum and Vaccine Research Institute.

Cell culture:

Vero cell line was obtained from the Virology Department of Animal Health Research Institute, Dokki, Giza, and used for propagation and titration of infectivity of the virus strains and in

RESULTS

The percentages of sera samples that contain antibodies to PPR and RP viruses and antibody titer for both viruses in 125 sheep from two breeds and 57 goats sera are shown in Tables, 1 & 2.

Table 1, showed that, Out of 68 sera samples from mixed breed sheep, 25(36.76%), 20 (29.41%), and 4 (5.88%) of the samples contained antibodies to PPR virus that ranged from 1:2 - 1:8, and one sample has a titer of 1:32. Out of 57 sera samples of Barki sheep, 20 (35.08%), 17 (29.82%), and

Table 1: Serum neutralizing (SN) antibodies for PPR virus in sheep and goats sera.

| Species & breeds | Nr. of samples. | SN antibody titer | | | | | | | | | |
|------------------|-----------------|-------------------|-------|-----|-------|-----|------|------|---|------|------|
| | | 1:2 | | 1:4 | | 1:8 | | 1:16 | | 1:32 | |
| | | nr. | % | nr. | % | nr. | % | nr. | % | nr. | % |
| Mixed sheep | 68 | 25 | 36.76 | 20 | 29.41 | 4 | 5.88 | 0 | 0 | 1 | 1.47 |
| Barki Sheep | 57 | 20 | 35.08 | 17 | 29.82 | 3 | 5.26 | 0 | 0 | 0 | 0 |
| Goats | 57 | 18 | 31.57 | 17 | 29.82 | 1 | 1.75 | 0 | 0 | 0 | 0 |

serum neutralization test (SNT). Growth medium was Eagle's minimal essential medium (MEM) supplemented with 10% newborn calf serum, 100 IU/ml penicillin and 100 ug/ml streptomycin. The maintenance medium was the same as growth medium except the serum concentration was reduced to 2%.

Serum neutralization test (SNT):

The infectivity of both PPR and RP viruses was titrated by the microtechnique according to Rossiter and Jesset, (1982) and the 50% end point of virus infectivity was determined according to Reed and Muench (1938). The sera were heat inactivated at 56C, for 30 minutes and the microtechnique of SNT was done according to Rossiter et al (1985) using 100 TCID₅₀ of both viruses.

(5.26%) have antibodies in a titer ranged from 1:2 - 1:8. Only. Examination of 57 goats sera revealed the presence of PPR antibodies in 18 (31.57%) (29.82%) samples in a titer ranged from 1:2 - 1:8 and only one sample (1.75%) has antibody titer of 1:8.

Table 2, indicated that, RP virus neutralizing antibodies are found in mixed breed sheep sera 8(11.76%), 2 (2.94%) and 5 (7.35%) samples with antibody titer ranged from 1:2 - 1:8, and one sample has a titer of 1:32. Barki sheep sera showed antibodies to RP virus in 10 (7.54%) 7 (12.28%), and 2(3.5%) out of the 57 samples examined and antibody titer ranged from 1:2-1:8, and only one sample (1.75%) has a titer of 1:16. Examination of 57 goats sera showed the presence of RP virus antibodies in 6(10.52%), 4 (7.01%), and 2 (3.5%) samples with a titer ranged from 1:2 - 1:8.

DISCUSSION

Examination of sera samples collected from two sheep breeds (Barki and Mixed breeds) as well as goats for the prevalence of the neutralizing antibodies for PPR and RP virus revealed that

Comparison of the percentages of the positive samples and the antibody titer for both viruses (Table, 1 & 2) revealed that the positive samples for PPR virus antibodies in Mixed sheep breed (29.41%), Barki sheep (29.82%) and goats

Table 2: Serum neutralizing (SN) antibodies for RP virus in sheep and goats sera.

| Species & breeds | Nr. of samples. | SN antibody titer | | | | | | | | | |
|------------------|-----------------|-------------------|-------|-----|-------|-----|------|------|------|------|------|
| | | 1:2 | | 1:4 | | 1:8 | | 1:16 | | 1:32 | |
| | | nr. | % | nr. | % | nr. | % | nr. | % | nr. | % |
| Mixed sheep | 68 | 8 | 11.76 | 2 | 2.94 | 5 | 7.35 | 0 | 0 | 1 | 1.47 |
| Barki Sheep | 57 | 10 | 17.54 | 7 | 12.28 | 2 | 3.5 | 1 | 1.75 | 0 | 0 |
| Goats | 57 | 6 | 10.52 | 1 | 7.01 | 2 | 3.5 | 0 | 0 | 0 | 0 |

(Table 1&2), at 1:4 serum dilution the percentage of sheep sera that contained antibodies to PPR virus was 29.41% in Mixed sheep and 29.82% in Barki sheep and the same rate (29.82%) in goats. This result indicated that there is no significant difference in the prevalence of PPR virus antibodies in the two sheep breeds examined and the prevalence of these antibodies is equally the same in sheep and goats. The antibody titer for PPR virus antibodies in sheep and goats ranged from 1:2-1:8 (Table 1), only in 1.47% of Mixed breed sheep the titer reached 1:32. In this respect, Taylor (1979) and Ekue et al. (1992) reported antibody titer for PPR virus in sheep ranged from 0.6-1.5 and in goats from 0.6-0.9. Obi et al. (1984) reported that sheep has PPR virus antibody titer higher than goats.

Examination of the same sera samples of sheep and goats for antibodies against RP virus (Table 2) revealed that at 1:4 serum dilution the prevalence of RP virus antibodies was higher in Barki sheep (12.28%) than in Mixed breed sheep (2.94%) and than goats (7.01%). In this respect, Zwart and Row (1966) found that 18.8% of sheep sera and 15.2% of goats contained antibodies to RP virus, Obi et al., (1984) reported higher prevalence of RP virus antibodies in sheep (19.5%) than in goats (6%).

(29.82%) were higher than that for rinderpest virus antibody positive samples which was, 2.94% 12.28% and 7.01% for the two sheep breeds and goats respectively, This result indicated that PPR virus neutralizing antibodies are more prevalent than that of RP virus in sheep and goats. This result conform that of Obi et al. (1984), Taylor (1979), Tawfik et al., (1989), Lefevre et al (1991) and Ekue et al (1992).

The results obtained in this study also agrees with that of Obi et al., (1984), Taylor and Abegunde (1979), in that there is some level of cross neutralization between PPR and RP virus antibodies and it is easy to distinguish the two viruses by titrating sera against both of them (Obi et al., 1964, Taylor, 1979).

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