

## RESPONSE OF PREGNANT SHEEP AND GOATS TO A SPECIFIC VACCINE FOR PESTE OF SMALL RUMINANTS

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Received: 14.5.2000.

Accepted: 27.8.2000.

### SUMMARY

Five susceptible pregnant dams of each of sheep and goats were vaccinated with PPR-attenuated vaccine, 50 to 60 days pre-partum. They, as well as, their offsprings remained clinically healthy. Post-partum serum PPR neutralizing antibody titres were generally lower than pre-partum peak titres in all animals. PPR antibody titres in the colostrum of these animals were generally appreciably higher than in their sera, throughout a test period of 48 hours post-partum. Reasonable serum titres of passively acquired PPR antibodies were fairly existing in the offsprings of both sheep and goats. Such antibodies persisted for 3 to 4 months with some individual variations. The majority of the offsprings reverted to seronegativity by the 5th month of age.

### INTRODUCTION

FAO (1996) stated that PPR was currently circulating with devastating effect in sheep and goats across the Asian continent from Arabia through Pakistan, India and as far as Nepal and Bangladesh. Animal disease status worldwide in 1997, as reported by OIE (1998) mentioned the occurrence of pest of small ruminants "PPR" in countries in western, central and eastern Africa, the middle east and south Asia. The disease was reported for the first time in that year in Pakistan and in Somalia. The disease had been reported to occur in Egypt in limited outbreaks (House, 1987; Ikram et al., 1988; Abo El-Hassan et al., 1989; El-Sanousi et al., 1989; Ismail and House, 1990; Ismail et al., 1990; El-Allawy et al., 1993 and Mouaz et al., 1995).

Before the development of a specific homologous PPR vaccine (Diallo et al., 1989), the

heterologous Plowright RBOK rinderpest vaccine was widely used to protect small ruminants against PPR (Bourdin et al., 1970; Bourdin, 1973; Taylor, 1979; Bonniwell, 1980; Adu and Nawathe, 1981; Abegunde, 1983; Bonniwell, 1983 and Obi, 1983).

A specific homologous PPR attenuated vaccine was developed in Egypt (Khodeir and Mouaz, 1998). It proved to be safe and immunogenic for sheep and goats through ample quality control criteria (Abeer, 1997 and Afaf, 1998). It was also evidenced that normal physiological functions of the vital organs were found to be maintained in vaccinated animals (Hanan, 1998). It was also proved to be of such same merits on being inoculated simultaneously with attenuated Rift valley fever virus vaccine (Mouaz et al., 1998) or with attenuated sheep pox vaccine (Samir et al., 1999), in synchronized vaccination trials in sheep. Hence, the object of the present work was aiming at experimenting such a locally produced PPR attenuated vaccine in pregnant sheep and goat to find out their response to vaccination as well as to envisage the serological status of their offsprings and the role of the colostrum in this respect.

## **MATERIALS AND METHODS**

### **MATERIALS :**

#### **Animals :**

Five pregnant dams of each of sheep and goats, seronegative to both PPR and rinderpest viruses

were used in the present study. Fifty to sixty days approximate before parturition, they were vaccinated with the attenuated PPR vaccine by dosing at  $10^3$  TCID<sub>50</sub>/ml/head, given S/C, in the region of the neck. Two other pregnant dams (one sheep and one goat) of the same condition were held as non vaccinated non contact control animals, kept in a separate pen. Test animals were closely observed and clinically inspected throughout the experimentation period. All animals as well as their offsprings were seromonitored for PPR neutralizing antibodies at predetermined indicated intervals. Colostra of all animals were also tested at targeted intervals.

#### **Serum samples :**

Serum samples were collected from all dams as well as from their offsprings at predetermined intervals throughout experimentation.

#### **Colostrum samples :**

Colostrum samples were collected by hand milking and were treated as the serum samples.

#### **Viruses :**

##### **Rinderpest virus:**

The cell culture attenuated rinderpest virus (RBOK) at its 99th passage on bovine kidney cells was used to produce rinderpest vaccine on VERO cells.

The vaccine virus was used to screen tested animals sera (Plowright et al., 1971).



## **Peste Des Petits Ruminants Virus (PPRV) :**

A VERO cell attenuated local strain (Egypt 87) of PPRV was used to screen tested animals sera and for vaccination as well as for quantifying PPR antibody titres in indicated test samples.

### **Cell cultures:**

Certified VERO cells were used in cultures for vaccine manufacture of either rinderpest or PPR as well as for virus titration and neutralization tests.

### **METHODS:**

#### **Virus titration:**

The infectivity titres of both rinderpest and PPR viruses were estimated according to the method described by Rossiter and Jessette (1982).

#### **Virus neutralization:**

Both quantitative and qualitative neutralization tests were done according to the method described by Rossiter and Jessette (1982). The neutralizing antibody titres were expressed as the reciprocal of the final serum or colostrum dilution inhibiting the CPE produced by 200 TCID<sub>50</sub>/0.1ml of PPR virus suspension.

## **RESULTS**

### **PPR vaccination of sheep and goat pregnant dams:**

#### **Clinical inspection:**

Vaccinated animals remained healthy and gave birth to healthy normal offsprings. Non of the dams aborted. No teratogenic features could be detected in all offsprings. There was no twining in all tested animals.

#### **Serological response to PPR vaccine:**

Fairly high PPR neutralizing antibody titres ranging between 64 and 256 were detected 3 and 4 weeks post vaccination of dams. Control animals remained seronegative. Post partum serum titres of mothers were dramatically of lower levels (32-128), throughout a 48 hours post partum test period.

#### **Colostrum PPR antibodies:**

Appreciably higher levels of PPR antibody titres could be estimated in colostrum samples of mothers, ranging between 64 and 256 throughout the first 17 hours post partum. Colostrum antibody titres ranging between 32 and 128 were found subsequently in samples collected at 24 and 48 hours post partum.

Table (1) : Serological response of pregnant sheep and goats to PPR vaccine and colostrum transfer of neutralizing antibodies to offsprings

Test of	Time of sampling	Animals No.	Sheep					Goats									
			1	2	3	4	5	6**	1	2	3	4	5	6**			
Pregnant dams	Just prior to vaccination 3 weeks post vaccination 4 weeks post vaccination	0*	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
		64	256	128	64	128	0	256	128	64	128	128	256	0	0	0	
		64	256	128	64	128	0	256	128	64	128	128	256	0	0	0	
Mothers	2 hours post partum 7 hours post partum 24 hours post partum 48 hours post partum	32	128	64	32	64	0	128	64	32	64	128	0	0	0	0	
		32	128	64	32	64	0	128	64	32	64	128	0	0	0	0	
		32	128	64	32	64	ND	128	64	32	64	128	0	0	0	0	
Colostrum	2 hours post partum 7 hours post partum 11 hours post partum 17 hours post partum 24 hours post partum 48 hours post partum	128	246	256	128	128	0	256	128	64	128	256	0	0	0	0	
		128	256	256	128	128	ND	256	128	64	128	256	0	0	0	0	
		128	256	128	128	128	0	256	128	64	64	128	256	ND	0	0	0
Offsprings	Just prior to suckling 2 days old 5 days old 10 days old 15 days old 30 days old 40 days old 60 days old 90 days old 120 days old 150 days old	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
		64	128	64	32	64	0	128	64	64	64	64	128	0	0	0	0
		64	128	64	32	64	ND	128	64	64	64	64	128	0	0	0	0
		64	128	64	32	64	0	128	64	32	64	64	128	ND	0	0	0
		64	128	64	32	64	ND	64	32	16	32	64	64	0	0	0	0
		64	128	64	16	32	0	64	32	16	32	64	64	0	0	0	0
		32	64	64	8	16	0	32	16	8	16	32	64	0	0	0	0
		16	64	64	4	8	0	32	8	4	8	32	64	0	0	0	0
		8	32	32	0	4	ND	16	4	0	4	16	32	0	0	0	0
		0	16	0	0	0	0	8	0	0	0	8	16	0	0	0	0

\* : PPR-neutralizing antibody titres expressed as the final dilution of serum or colostrum inhibiting the CPE of PPR virus (200 TCID<sub>50</sub>/0.1) on Vero cells.  
 \*\* : Non vaccinated non contact control animals.  
 ND : Not Done.



Table (2): Geometric mean titres of PPRV-antibody in pregnant sheep, goats, mothers, colostrum and offsprings.

Test of	Time of sampling	Animals	
		Sheep "5"	Goats "5"
Pregnant dams	Just prior to vaccination	0*	0
	3 weeks post vaccination	111.4	147
	4 weeks post vaccination	111.4	147
Mothers	2 hours post partum	55.7	73.5
	7 hours post partum	55.7	73.5
	24 hours post partum	55.7	73.5
	48 hours post partum	55.7	73.5
Colostrum	2 hours post partum	168.9	147
	7 hours post partum	168.9	147
	11 hours post partum	147	147
	17 hours post partum	128	128
	24 hours post partum	84.4	84.4
	48 hours post partum	73.5	73.5
Offsprings	Just prior to suckling	0	0
	2 days old	64	84.4
	5 days old	64	84.4
	10 days old	64	73.5
	15 days old	64	73.5
	30 days old	48.5	36.8
	40 days old	48.5	36.8
	60 days old	27.9	18.4
	90 days old	16	12.1
	120 days old	11.3	8
150 days old	16	8	

\* PPR- neutralizing antibody titres were expressed as the reciprocal of the final dilution of serum or colostrum inhibiting the CPE of PPR virus (200 TCID<sub>50</sub>/0.1ml) on Vero cells.

Fig. (1): Geometric mean titres of PPRV- antibody in pregnant sheep, goat, mothers and colostrum.

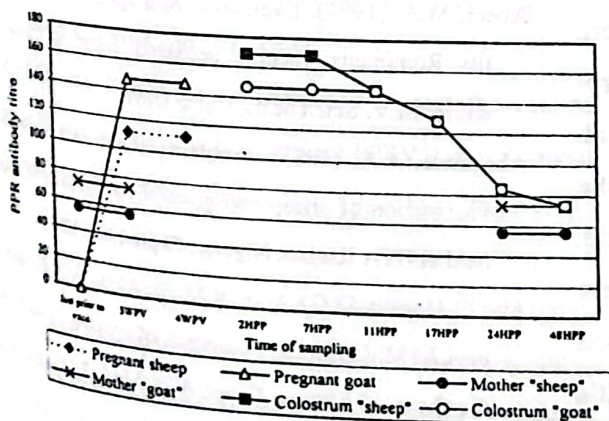
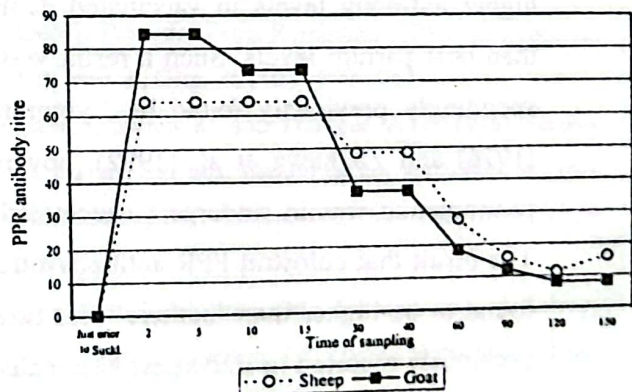


Fig. (2): Geometric mean titres of PPRV- antibody in sheep and goat offsprings.



Colostrum transfer of PPR antibodies to offsprings: Offsprings were seronegative before suckling the colostrum. Seroconversion occurred after nursing. PPR neutralizing antibody titres ranging between 16 and 128 were traced up to the 40th day of age. At the age of 120 days, two sheep showed titres of 16 and 32. At the same age, two goats maintained a titre of 16. By the 5th month of age, only one sheep was seropositive in a titre of 16. Two goats of a serum titre of 8 were found at the same age.

## DISCUSSION

Although the number of sheep and goats used in this study was only five of each, PPR vaccine seemed to be reasonably well tolerated by heavily pregnant animals. Negativity of obvious teratogenicity and absence of abortion would be considered additional proof of safety of this vaccine. Serological response in terms of neutralizing PPR antibody was fairly of appreciable levels in vaccinated dams, mothers, colostrum and in nursing offsprings. Tabulated results in this study denoted higher antibody levels in vaccinated dams' sera than post partum levels. Such a result was correspondingly previously found by Lyigoren et al. (1976) and Zaghawa et al. (1992), however the phenomenon was in rinderpest vaccinated cattle. The result that colostrum PPR antibody titres were found to be higher than mothers' sera titres, was previously reported in rinderpest vaccinated cattle by Brown (1958). Our results seemingly indicate

that the titres and duration of colostrally transferred PPR antibody in newborn sheep and goats incline to be directly proportional to the maximum titre acquired after nursing which in turn depends on the level of colostrum antibody received within the neonatal period.

It seems that natural passively acquired PPR antibodies decline gradually at a uniform rate till disappear completely by approximately the fifth month of age. Correspondence in this respect was previously reported with rinderpest in cattle (Brown, 1958) and polio in infants (Martins et al., 1958). It could be withdrawn that PPR attenuated vaccine is safe and immunogenic for pregnant sheep and goat dams. It has no deleterious effect on them neither on their offsprings. In addition, newborn sheep and goats suckling the colostrum enjoy the acquisition of passively transferred PPR antibody that might be persisting up to the 3rd or 4th month of age.

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