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Blood Polymorphism in West African Breeds of Sheep

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ABSTRACT

This paper reports the blood groups and blood protein distribution in West African sheep breeds. About 100 animals of the Djallonke, Fulani and Touabire breeds were sampled for blood polymorphism analysis. Their blood groups were typed by haemolytic and agglutination reactions, and their blood proteins by starch gel electrophoresis. Almost all the loci analysed showed variability in the three breeds, with the Touabire and Fulani being closer to each other than to the Djallonke.

Keywords: blood groups, breed, haemoglobin, sheep, serum protein, transferrin

INTRODUCTION

In West Africa, two types of hairy thin-tailed sheep have been reported: the savannah and the forest types (Epstein, 1971). The savannah type in the tsetse fly-free area is large, long-legged and trypanosusceptible, while the dwarf forest type found in the humid zone is trypanoresistant. Bradley (1995) demonstrated an increasing introgression of trypanosusceptible blood into trypanoresistant cattle as a consequence of the increase in transhumance in recent decades due to drought and to farmers' preference for large animals.

Since genetic diversity is a basic requirement for animal improvement, it needs to be preserved through characterization and management. Blood groups and blood proteins have been used widely to characterize animal populations because of their polymorphism and their simple mode of inheritance. Moreover, they are supposed to be linked to productivity traits and environmental adaptation (Dally *et al.*, 1980; Vicovan and Rascu, 1989; Charon *et al.*, 1996). However, data on blood polymorphism in tropical African sheep are scarce and refer only to their haemoglobin types (Olusanya, 1975; Ndamukong, 1995).

This paper reports the distribution of blood groups and blood proteins in three West African breeds of sheep.

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MATERIALS AND METHODS

Animals

Blood samples were collected from Senegalese Djallonke (dwarf forest type). Peul-peul (Fulani) and Touabire (Savannah type) breeds of sheep. The Fulani sheep were sampled in the Djoloff, located 250 km from Dakar in north Senegal, while the Djallonke animals were chosen in Kolda, south Senegal, about 500 km from the capital. The Touabire sheep, which originated from Mauritania, were sampled in Dakar market, where they are brought for sale.

Blood samples (5 ml) were collect ed from approximately 100 animals per breed. The animals were adults of both sexes and were chosen from different flocks. The blood samples were collected into sodium citrate and sent to the Laboratoire d'Analyses Génétiques pour les Espèces Animales (LABOGENA), Jouy-en-Josas. France, for analysis.

Blood typing and data analysis

The blood protein polymorphism was analysed by starch gel electrophoresis and by electrofocalisation in the case of haemoglobin (Nguyen and Bunch, 1980).

Six OEA (ovine erythrocyte antigen) systems were analysed by haemolytic and haemagglutination reactions as described by Nguyen (1972). Blood protein alleles were determined by direct counting; blood group 'allele' frequencies were calculated after Hardy-Weinberg equilibrium had been established at the transferrin and carbonic anhydrase loci.

RESULTS

Blood group polymorphism

The distribution of blood groups in the three sheep breeds is shown in Table 1. The observed allelic frequencies may be biased because of the low number of animals used. However, according to Nei (1978), the error is low in sample sizes of 100. Except for locus B, which was very polymorphic, all the other alleles were present in the three breeds at different frequencies. The differences between the Djallonke and Fulani sheep were statistically significant (p < 0.05) for alleles A^{ab} , C^{ab} , D^{a} and **R**, the Touabire sheep being intermediate.

Bloodproteins

Systems	Allelles	Frequencies		
		Djallonke	Fulani	Touabire
A	a	0. 526	0. 55	0. 499
	b	0. 087	0. 035	0.068
	a b	0. 025	0.001	0. 021
В	b	0. 03	0. 181	0. 161
	a b	0.005	0.027	0. 108
	abe	0	0. 022	0
	abi	0	0. 023	0
	hi	0. 020	0.051	0. 148
	d	0. 05	0.007	0. 027
	di	0	0	0.046
	е	0. 05	0.014	0. 013
	ei	0	0	0. 038
	i	0. 025	0. 079	0. 020
	а	0	0. 022	0
	а	0. 005	0.045	0.071
	ab	0.005	0.056	0.063
D	а	0. 529	0. 341	0. 471
М	a	1	0. 99	0. 975
R	r	0. 326	0. 44	0. 408
Accordance	with Hardy-Weinberg la	aw"		
Tf: χ^2 : 4 df	. 0	4.64	3. 25	4.92
CA: χ^2 : ldf		0. 038	0.756	0. 277

TABLE I OEA allele frequencies in West African breeds of sheep

^aTf, transferrin; CA, carbonic anhydrase: df, degrees of freedom

Djallonke sheep in Cameroon (Ndamukong, 1995) but contradicts results obtained in the same breed in Nigeria by Olusanya (1975). Concerning transferrin, seven phenotypes (AA, CC, AD, AB, AC, AD and CD) were observed, the gene alleles being TfA, TfB, TfC and TfD. The pattern of distribution of the B and D alleles (respectively low and high in the Djallonke breed and medium in the other breeds) confirmed the results obtained **in** trypanosusceptible and trypanorasistant cattle by

	Alleles	Allele frequencies		
Systems		Djallonke	Fulani	Touabire
Transferrin	A G B C D	0.216 0 0.05 0.110 0.609	0.428 0.041 0.072 0.139 0.320	0.308 0.030 0.142 0.126 0.394
Haemoglobin	$egin{array}{c} A \ B \end{array}$	0 1	0 1	0.15 0.985
Carbonic anhydrase	M S	0.232 0.768	0.08 0.920	0.05 0.95
Protein X	Х	0.112	0.094	0.089

TABLE II Distribution of blood protein in West African breeds of sheep

DISCUSSION

Although several sheep breeds have been described in West Africa (Doutressole, 1947; Epstein, 197 1), little is known about the extent of genetic diversity between and within these breeds (Rege, 1994). In this paper, we analyse the value of blood groups and blood protein polymorphism in the genetic study of sheep breeds. The overall results showed significant differences between the Djallonke breed on the one hand and the Touabire and Fulani breeds on the other. In a spatial autocorrelation study, Ordas and Carriedo (1996) showed that the allele frequency differences among European sheep breeds were due to migration and genetic drift. This could support the existence of differences in the historic expansion into West Africa of Djallonke and the other breeds and is consistent with the classification of Epstein (1971). In fact, this author pooled the Touabire and Fulani breeds in the same group (the Savannah) while the Djallonke breed was classified as a forest type.

CONCLUSION

Analysis of the biochemical polymorphism in West African sheep breeds showed differences among breeds which reflected their historic expansion. However, this work needs to be continued on a greater number of breeds and with other markers (microsoftallite). The relationship bet waan class proteins and economically relevant

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Polimorfismo sanguineo en razas ovinas de Africa Occidental

Resumen – Este artículo estudia la distribución de proteinas plasmáticas y grupos sanguineos en razas ovinas de Africa Occidental. Se obtuvieron muestras de sangre de alrededor de 100 animales de las razas Djallonke, Fulani y Touabire, con objeto de estudiar su polimorfismo sanguineo. Los grupos sanguineos se tipificaron mediante reacciones de hemólisis y aglutinación, y las proteinas sanguineas mediante electroforesis en gel. Casi todos los *loci* analizados mostraron variabilidad en las 3 razas, siendo las razas Touabire y Fulani más parecidas entre si que a la raza Djallonke.