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Electrophoresis as an Aid in Detecting Pathological Conditions in Wild Mammals¹

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ABSTRACT

Changes in the electrophoretic distribution of plasma proteins were noted in wild animals with a variety of parasitic and infectious diseases. This technique may be useful in screening for disease before death or sacrifice of animals.

INTRODUCTION

Alterations in the electrophoretic distributions of plasma and serum proteins have been described in various animals both in spontaneous diseases^{2 3 5} and in the screening of experimental animals¹. The alterations in normal plasma protein distribution following disease are largely nonspecific consisting of an increase in the globulin fractions and a decrease in the albumin component.

During studies of the plasma protein patterns of wild mammals of the USAEC Oak Ridge Reservation, electrophoretic patterns of some diseased mammals differed markedly from the normal distribution for the particular species. Further studies were carried out to determine the disease present, the plasma protein pattern, and the possible relationship between the two.

MATERIALS AND METHODS

Blood coagulation was prevented by EDTA, and 6-12 µl samples of plasma were analyzed for proteins using the Spinco Model R paper electrophoresis system, staining with brom-

phenol blue, and scan-tracing the protein fractions with a Spinco Model R Analytrol. Calculations of protein fractions and albuminto-globulin ratios (A/G) were made from the scan-tracing of each sample.

RESULTS AND DISCUSSION

The effects of several pathological conditions upon plasma protein distribution are summarized in Table 1. With the exception of the fox with uncomplicated rabies, all of the diseased animals showed a marked depression in the A/G. This is the result of concurrent albumin decrease and variable increases in alpha, beta1, and beta2 globulins. In myiasis and mite infestations of rabbits and skunks, where the number of parasites was high, inflammation with serous exudate was observed and involved approximately two to twenty percent of the hosts' total body surface area. Increased vascular permeability associated with the inflammatory process may have resulted in some loss of albumin from the circulation. Inflammation was associated with the response to Cuterebra infestation in chip-

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munks and white-footed mice where purulent exudate and/or abscess formation was observed. The presence of pneumonia with the associated extensive tissue involvement in one rabid fox may account for the depression in A/G in this animal when compared to the one with rabies alone.

Since variations in plasma protein patterns appear to accompany many abnor-

mal conditions, the use of this technique in experimental and wild animals is worthwhile. The small size of the blood serum sample (.006 - .012 ml) required and the relative simplicity of the procedure make it suitable for establishing basic blood parameters and detecting responses of the host to injurious agents in a wide range of pathological condi-

Table 1. Percent plasma protein distribution in wild mammals with and without disease.

Species	Commor	name	Number ani	mals Dise	ase		
				Globulins			
Album	in	a 1	a ₂	$oldsymbol{eta_1}$	β2	γ	A/G
Oryctolag	ue cunicu	lus (dor	nestic rabbit),	five animals	no disease.		
61.8±1.		2±0.21	5.8±0.25	8.8±0.60		16.4±1.31	1.640±0.122
O. cunicu	lus, five	animals,	caliphorid my	iasis and ac	arine mange.		
53.4±0.		±0.39	6.7±0.45	8.6±0.54		23.3±0.70	1.154±0.057
Tamias si	triatus (ea	stern ch	ipmunk), seve	n animals, r	o disease.		
23.6±1.	.06 9.5	± 1.69	18.8±2.48	8.2±0.71	15.1±3.05	10.0 ± 2.68	0.656±0.098
T. striatu	s, two an	imals, m	yiasis, Cutebra	r sp.			
13.8±2.	95 12.9	±3.15	16.9±1.80	10.3 ± 2.7	20.5 ± 1.10	17.5±5.70	0.284±0.075
Peromysca	us leucopi	us (whit	e-footed mous	e) fifteen a	inimals, no dise	ase.	
47.4±1.		±0.7	10.7 ± 1.0	21.3 ± 0.6		11.8 ± 0.8	0.899±0.034
			lls, myiasis, Cr	iterebra ang	ustifrons.**		
$32.9 \pm 1.$		±0.5	15.6±1.0	32.0±1.5		11.3 ± 0.9	0.490 ± 0.032
Sigmodon	hispidus	, (Cotto	n rat) two an				
$43.9 \pm 0.$		±1.03		13.8±0.62		19.9 ± 0.96	0.805 ± 0.001
			ilariasis, <i>Litom</i>				
36.9±1.		±0.85		19.2±1.72		16.8 ± 0.40	0.585±0.035
			(gray fox), or				
37.4	5.0		6.8	13.9	22.9	14.0	0.597
			nimal, rabies.			_	
37.5	13.6		7.8	28.0	6.5	6.6	0.600
			mal, rabies an				
30.2	9.4		5.7	20.8	25.3	8.6	0.433
			l skunk), five			120.00	0.550 . 0.555
	04 16.7			13.3±0.02	17.7±0.008	13.2 ± 0.02	0.658 ± 0.107
			carine mange.		an .		
22.2	16.7			20.4	27.6	13.1	0.285

^{*} Mean ± standard error of the mean.

** Data from Payne et al.4

NOTE: Pre-albumin determinations for normal chipmunks were 14.8±3.04, and for diseased chipmunks, 8.1 ± 0.937 .

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