

HEMATOLOGY AND OCCURRENCE OF HEMOPARASITES IN MIGRATING SHARP-SHINNED HAWKS (*Accipiter striatus*) DURING FALL MIGRATION

LAUREN V. POWERS AND MARK POKRAS

Tufts Wildlife Clinic, Tufts University School of Veterinary Medicine, 200 Westboro Road,
North Grafton, MA 01536 U.S.A.

KIM RIO

304 Hellertown St., Hellertown, PA 18055 U.S.A.

CATHY VIVERETTE AND LAURIE GOODRICH

Hawk Mountain Sanctuary, RD 2, Box 191, Kempton, PA 19529 U.S.A.

ABSTRACT.—Packed cell volume (%), total solids (g/dl), white blood cell count (cells/ μ l), differential and absolute white blood cell counts, and prevalence of hemoparasites were determined for 85 healthy sharp-shinned hawks (*Accipiter striatus*) during the 1991 fall migration. The packed cell volume ($47.6 \pm 6.73\%$), total solids (2.83 ± 0.58 g/dl) and white blood cell count ($12\,900 \pm 7\,310$ cells/ μ l) are within ranges reported previously for other raptors, both captive and wild. Immature birds showed a greater prevalence for the hemoparasites *Hemoproteus* and *Leukocytozoon* than adults but there was no significant difference in prevalence between males and females. These findings add to the small but growing data base on hematology of birds of prey.

KEY WORDS: hematology; hemoparasites; migration; sharp-shinned hawk; stress.

Hematología y ocurrencia de hemoparásitos en *Accipiter striatus* migratorios durante la migración otoñal de 1991

RESUMEN.—Se determinó el volumen celular (%), sólidos totales (g/dl), conteo de glóbulos blancos (células/ μ l), conteo diferencial y absoluto de glóbulos blancos y prevalencia de hemoparásitos para 85 individuos saludables de *A. striatus* durante la migración otoñal de 1991. El volumen celular ($47.6 \pm 6.73\%$), los sólidos totales (2.83 ± 0.58 g/dl) y el conteo de glóbulos blancos ($12\,900 \pm 7\,310$ células/ μ l), están dentro de los rangos reportados previamente para otros rapaces, tanto cautivos como silvestres. Aves inmaduras mostraron una mayor prevalencia de los hemoparásitos *Hemoproteus* y *Leukocytozoon* que los adultos, pero no hubo una diferencia significativa en la prevalencia entre machos y hembras. Estos resultados se agregan a la pequeña pero creciente base de datos sobre la hematología de aves de presa.

[Traducción de Ivan Lazo]

Hematology is a useful tool in determining normal and pathological states in a variety of species including birds. The number of studies on raptor hematology is limited despite the large amount of information available for poultry. In addition, most reports are restricted to captive individuals or birds under rehabilitation and have a small sample size. Previous studies (Hunter and Powers 1980, Gessaman et al. 1986) have examined hematology from free-ranging raptors, which may vary from values obtained from captive birds. The purposes of this study were to add to the data base for hematology of wild raptors and to examine hematologic data for evidence of systemic disease.

MATERIALS AND METHODS

Subjects of our study were migrating sharp-shinned hawks (*Accipiter striatus*) captured for banding purposes at the Little Gap Banding Station near Hawk Mountain Sanctuary in eastern Pennsylvania. Birds were trapped from 9 September to 11 November 1991 between 0630 and 1635 H. Captured birds were weighed to the nearest gram. Each bird was classified as either immature (hatching year) or adult (second year or older) based on plumage characteristics.

Approximately 1 ml blood was collected from the jugular vein. Captured birds were injected with approximately 1 ml lactated Ringer's solution subcutaneously before release. A drop of whole blood was smeared onto a glass slide for measurement of the differential white blood cell count. A sample of whole blood was placed in a mi-

rohematocrit tube (approximately 70 μ l) and spun at 10000 rpm for 4 min to measure packed cell volume (PCV), or hematocrit. Total solids (TS), an estimate of the protein content, was measured from the plasma using a clinical refractometer.

Samples of whole blood were diluted and stained using a Unopette[®] Eosinophil staining kit (Becton-Dickinson and Co., Test No. 5877). Stained granulocytes (heterophils, eosinophils and basophils) were counted manually using a hemacytometer. Total white blood cell count (WBC) was calculated using methods described by Dein (1984): cells in 10 squares were counted and the result multiplied by 32 to adjust for dilution. This value was divided by the percentage of staining granulocytes in the differential white blood cell count to adjust for the lack of staining of lymphocytes and monocytes. Blood smears were dipped in the quick stain HEMA-3[®] (Biochemical Sciences, Inc.) for differential white blood cell counting. One hundred leukocytes were counted and the percent of each cell type was recorded. Differential counts were verified by having each slide read by three different analysts. Absolute white cell counts for each of the types of leukocyte were calculated by multiplying total white blood cell count by the percentage of each cell type in the differential count. Smears were also examined for the presence of blood parasites.

Mean, standard deviation, range, median, and percentiles are reported for the hematologic data. Analysis of variance (ANOVA) and Student *t*-tests were used to detect significant differences in selected parameters. An alpha level of less than 0.05 was considered significant. The Statistical Analysis System (SAS 1988) was used for analysis of data.

RESULTS

Results for packed cell volume (%), total solids (g/dl), white blood cell count (cells/ μ l), relative differential white blood cell count (% cell type), and absolute differential count (cells/ μ l) are presented in Table 1. We found that distributions of PCV, WBC, relative differential counts for heterophils, basophils, and monocytes, and absolute differential counts for all leukocyte types were not statistically normal (Kolmogorov statistic, *W*: Normal <95%). Therefore, percentiles may be more appropriate for accurate reporting of these data. Comparisons between males and females and between adults and immature birds found no significant differences in hematologic parameters, so results from all birds are compiled in Table 1. Prevalences (% affected) of the blood parasites *Hemoproteus* and *Leukocytozoon* detected from the smears are reported in Table 2. Comparison between males and females found no significant difference for either parasite. Comparison between adults and immature birds found that immature birds had a higher prevalence than adults for both *Hemoproteus* and *Leukocytozoon*. The hemo-

Table 1. Hematology of sharp-shinned hawks captured at Hawk Mountain Sanctuary.

N	PCV (%)	TS (g/dl)	WBC (cells/ μ l)	RELATIVE DIFFERENTIAL WBC COUNT (%)				ABSOLUTE DIFFERENTIAL WBC COUNT (cells/ μ l)				
				HET ^a	EO-SIN ^b	BASO ^c	LYMPH ^d	MONO ^e	HET ^a	EOSIN ^b	BASO ^c	LYMPH ^d
Mean	47.6	2.83	12900	81	81	81	81	72	72	72	72	72
SD	6.73	0.58	7310	27.0	7.67	0.40	63.4	3200	955	46	8480	187
Range	30-76	1.4-5.0	2400-37600	14.2	4.18	0.79	14.4	2800	715	110	5350	296
Median	47	2.8	11520	5-69	0-18	0-4	28-90	240-18800	0-3380	0-739	1770-23700	0-1640
50% percentile	44-52	2.4-3.2	7680-16800	26	7	0	65	2470	862	0	7930	111
95% percentile	38-56	2.0-3.8	4240-25400	16-34	5-11	0-1	54-75	1810-3630	422-1270	0-22	4030-11800	0-224
				8-52	1-15	0-2	35-83	820-9080	80-2520	0-241	1980-17500	0-744

^a Heterophils.
^b Eosinophils.
^c Basophils.
^d Lymphocytes.
^e Monocytes.

SITES

tial and sharp-6.73%), ranges valence ference ase on

otoñal

ilulas/viduos %), los dentro iduras lultos, dos se

Lazo]

arp-shinned ng purposes k Mountain ere trapped tween 0630 the nearest ture (hatch-on plumage

om the jug- h approxi- neously be- ared onto a white blood ed in a mi-

Table 2. Comparison of hemoparasite prevalence between immature and adult sharp-shinned hawks.

	TOTAL (N = 83)	IMMATURE (N = 60)	vs. ADULT (N = 23)	t	df	P
<i>Hemoproteus</i>	20.5%	28.3%	0.00%	2.98	81	0.0038 ^a
<i>Leukocytozoon</i>	16.9%	21.7%	4.35%	2.51	75	0.0143 ^a

^a Student's *t*-test. Probability <0.05 considered significant.

parasite *Plasmodium* was not detected on the blood smears.

Birds with white blood cell counts falling in the second and third quartiles are considered normal for this study. Those with a count above the third quartile are considered leukocytic. Those with a count below the first quartile are termed leukopenic. Comparisons of relative and absolute differential white blood cell counts between normal and leukocytic birds are presented in Table 3. Leukocytic birds had a significant absolute heterophilia, eosinophilia and lymphocytosis. However, differences were not significant between the two groups in relative differential white blood cell counts. Comparisons of relative and absolute differential white blood cell counts between normal and leukopenic birds are presented in Table 4. Leukopenic birds had a significant absolute heteropenia, eosinopenia, and lymphopenia and a significant relative heterophilia.

DISCUSSION

The mean hematocrit value of 47.6% falls within the range of reported values for trapped sharp-

shinned hawks ($49.5 \pm 2.5\%$) given by Gessaman et al. (1986) and agrees with previously reported values for other falconiforms (Bond and Gilbert 1958, Elliott et al. 1974, Cooper 1975, Balasch et al. 1976, Smith and Bush 1978, Hunter and Powers 1980, Gee et al. 1981, Ferrer et al. 1987). Hematocrits of migrating, healthy birds of prey may be significantly greater than those in captivity. High altitude is known to increase the production of red blood cells due to the greater demand for efficient oxygen extraction from air. Hemoconcentration from dehydration may occur in migrating birds (Carpenter 1975, Gessaman et al. 1986, Perry et al. 1986) because of relative reduced intake of water. No significant difference was found between the hematocrits of males and females, although it has been suggested that male birds should have a higher hematocrit than females due to the erythropoietic effect of androgens (Gee et al. 1981, Sturkie 1986). Gessaman et al. (1986) found no significant difference in hematocrits between the sexes of trapped sharp-shinned hawks. Hunter and Powers (1980) and Snyder et al. (1980) also found no significant difference between the sexes in either

Table 3. Comparison of differential hematology between sharp-shinned hawks with white blood cell counts above the third quartile (leukocytic) and falling in the second and third quartile (normal).

	TOTAL (N = 81)	LEUKOCYTIC (N = 18)	vs. NORMAL (N = 37)	t	df	P
RELATIVE COUNTS (%)						
Heterophils	27.0	22.1	25.1	0.79	53.0	0.430
Eosinophils	7.67	6.44	8.54	1.82	53.0	0.074
Basophils	0.40	0.28	0.38	0.57	53.0	0.574
Lymphocytes	63.4	69.5	64.4	-1.37	53.0	0.176
Monocytes	1.60	1.61	1.57	-0.07	53.0	0.947
ABSOLUTE COUNTS (cells/ml)						
Heterophils	3200	5270	2880	-2.22	19.1	0.038 ^a
Eosinophils	955	1420	1000	-2.11	53.0	0.039 ^a
Basophils	46	72	43	-0.66	20.0	0.518
Lymphocytes	8480	15400	7720	-6.88	24.4	0.0001 ^a
Monocytes	187	321	163	-1.29	19.8	0.21

^a Student's *t*-test. Probability <0.05 considered significant.

Table 4. Comparison of differential hematology between sharp-shinned hawks with white blood cell counts below the third quartile (leukopenic) and falling in the second and third quartile (normal).

	TOTAL	LEUKOPENIC	vs.	NORMAL	t	df	P
RELATIVE COUNTS (%)	(N = 81)	(N = 26)		(N = 37)			
Heterophils	27.0	33.0		25.1	-2.27	61.0	0.027 ^a
Eosinophils	7.67	7.27		8.54	1.13	61.0	0.263
Basophils	0.40	0.50		0.38	-0.52	37.5	0.607
Lymphocytes	63.4	57.8		64.4	1.90	61.0	0.063
Monocytes	1.60	1.65		1.57	-0.20	60.7	0.842
ABSOLUTE COUNTS (cells/ml)	(N = 72)	(N = 17)		(N = 37)			
Heterophils	3200	1720		2880	-3.39	48.7	0.001 ^a
Eosinophils	955	361		1000	-5.24	51.9	0.0001 ^a
Basophils	46	26		43	-0.75	52.0	0.454
Lymphocytes	8480	2840		7720	-9.88	46.7	0.0001 ^a
Monocytes	187	98		163	-1.72	50.1	0.091

^a Student's t-test. Probability <0.05 considered significant.

trapped or captive American kestrels (*Falco sparverius*). Age may also have an effect on hematocrit in that immature birds should have higher hematocrit values than adults (Rehder et al. 1982). No significant difference was found between immature birds and adults for hematocrit values in this study or that by Gessaman et al. (1986).

The mean total solids value of 2.83 g/dl was less than that reported previously for some falconiforms (Elliott et al. 1974, Halliwell et al. 1975, Smith and Bush 1978, Gee et al. 1981, Ferrer et al. 1987) but within the range reported by Balasch et al. (1976) and Snyder et al. (1980). Accurate comparisons with previously reported results may not be possible due to the different techniques in measuring protein levels in blood such as the Biuret and refractometric methods. Lumeij and de Bruijne (1985) showed that total solids measured with a refractometer have little correlation with serum protein levels measured by the Biuret method in rock doves (*Columba livia*). A low total solids value is often indicative of poor nutrition in raptors (Smith and Bush 1978, Ferrer et al. 1987). No significant difference in total solids was found between males and females or between adults and immature birds. These findings agree with those reported previously for some falconiforms (Snyder et al. 1980).

The prevalence of *Hemoproteus* in immature sharp-shinned hawks was 28.3%, yet no adults showed *Hemoproteus* parasitemia on the blood smears. On a similar note, 21.7% of immature birds had *Leukocytozoon* but only 4.35% of adults showed signs of

infection. No significant difference in hemoparasite prevalence was found between males and females. This finding is in agreement with previous reports (Kirkpatrick and Suthers 1988, Davidar and Morton 1993). The greater prevalence of *Hemoproteus* and *Leukocytozoon* in immature sharp-shinned hawks from this study is in contrast to other reports. Kirkpatrick and Suthers (1988) found that hatching-year birds were infected at a lower rate than older birds representing 59 species from central New Jersey. Yearling purple martins (*Progne subis*) were significantly less infected with *Hemoproteus prognei* than adults (Davidar and Morton 1993). Ashford et al. (1990) reported a higher prevalence of *Leukocytozoon toddi* in adult sparrowhawks (*Accipiter nisus*). This study also suggested a vertical mode of transmission from adults to young in the nest through vector species (*Culicoides* for *Hemoproteus* and ornithophilic members of the family Simuliidae for *Leukocytozoon*). Relapses of hemoparasitemia as birds become stressed or begin breeding has been suggested (Peirce 1980). A loss of detectable levels of hemoparasite in the peripheral blood (latency) between the months of October and April with a spring relapse was reported by Ashford et al. (1990). It is not unreasonable to suggest that immature sharp-shinned hawks become infected in the nest and are less able to achieve latency of infection with hemoparasites due to their naive immunologic status and because of the stresses of first migration and incompletely developed hunting skills. The clinical effects of hemoparasites have not been completely deter-

Table 5. Total and relative differential white blood cell counts reported in various raptors.

	N	WBC (cells/ $\mu\text{l} \times 10^3$)	RELATIVE DIFFERENTIAL WBC COUNT (%)						AUTHOR
			HETEROPHILS	EOSINOPHILS	BASOPHILS	LYMPHOCYTES	MONOCYTES		
Common buzzard (<i>Buteo buteo</i>)	11	14.0-49.0	20.5-39.8	5.5-19	0.25-8	35-65.5	0.25-3.75	a	
Honey buzzard (<i>Pernis apivorus</i>)	1	10.5	29.8	9	4.8	55.3	1.25	a	
Andean condor (<i>Vultur gryphus</i>)	1	13.5	42.8	11	2.5	42	1.8	a	
Golden eagle (<i>Aquila chrysaetos</i>)	1	23.5	52.5	6	3.3	34	4.3	a	
White-tailed sea eagle (<i>Haliaeetus albicilla</i>)	1	19.5	32.3	9.5	1.3	55	2	a	
Imperial eagle (<i>Aquila heliaca</i>)	1	15.0	44	11.8	2.3	40.3	1.3	a	
Tawny eagle (<i>Aquila rapax</i>)	1	42.5	57.3	10	2	30	0.8	a	
Marsh harrier (<i>Circus aeruginosus</i>)	3	9.0-33.0	26.5-39.5	1.5-6.5	2.8-5.3	48-59.5	2.5-10.5	a	
Common kestrel (<i>Falco tinnunculus</i>)	5	14.5-57.0	11.3-33	8.75-59.3	1.5-3.8	24-57.5	0.25-3.0	a	
Black kite (<i>Milvus migrans</i>)	5	10.0-28.0	28.8-35.3	12.8-35.5	2.3-3.5	29.5-50.5	0-2	a	
Red kite (<i>Milvus milvus</i>)	1	12.0	19.5	28.3	2.8	48.8	0.75	a	
Egyptian vulture (<i>Neophron percnopterus</i>)	1	29.5	43.8	5.5	8.5	37.5	4.8	a	

Table 5. Continued.

	N	WBC (cells/ $\mu\text{l} \times 10^3$)	RELATIVE DIFFERENTIAL WBC COUNT (%)					AUTHOR
			HETEROPHILS	EOSINOPHILS	BASOPHILS	LYMPHOCYTES	MONOCYTES	
King vulture (<i>Sarcorampus papa</i>)	2	41.9						b
Hooded vulture (<i>Necrosytes monachus</i>)	1	22.4						b
Savannah hawk (<i>Heterospizias meridionalis</i>)	1	31.0						b
Ornate hawk-eagle (<i>Spizaetus ornatus</i>)	1	33.0						b
White-bellied sea eagle (<i>Haliaeetus leucogaster</i>)	1	22.0						b
Crested serpent eagle (<i>Spilornis cheela</i>)	1	22.0						b
Bald eagle (<i>Haliaeetus leucocephalus</i>)	1	11.3						b
Collared forest falcon (<i>Micrastur semitorquatus</i>)	1	26.4						b
Crested caracara (<i>Polyborus plancus</i>)	2	24.1						b
Red-tailed hawk (<i>Buteo jamaicensis</i>)	2	6-46						c
Harris' hawk (<i>Parabuteo unicinctus</i>)	2	12-14						c

^a Christoph and Borowski (1961).

^b Elliott et al. (1974).

^c Halliwell et al. (1975).

mined. Hemoparasites have been shown to increase rehabilitation time in raptors (Olsen and Gaunt 1985). However, most studies on wild birds show no evidence of decreased longevity or reproductive ability in infected birds (Kirkpatrick and Suthers 1988, Ashford et al. 1990, Davidar and Morton 1993), although a negative effect on mate selection has been suggested (Kirkpatrick and Suthers 1988, Ashford et al. 1990, Davidar and Morton 1993). No significant differences in measured hematologic parameters were identified between hemoparasitemic and non-hemoparasitemic birds in our study, indicating no correlation between infection and other hemogram indicators of general health status.

The mean total WBC was $12\,900 \pm 7\,310$ cells/ μ l. This value falls within range of normal counts for other falconiforms (Elliott et al. 1974, Halliwell et al. 1975, Smith and Bush 1978). The relative differential white cell counts also agree with those previously reported (Table 5). It is difficult, if not impossible, to measure normal white blood cell counts accurately in raptors since these birds are unavoidably stressed when handled and especially when captured in the wild. Migration is also a source of stress to a raptor. ACTH and corticosteroids have been shown to be elevated during periods of stress in birds (Wolford and Ringer 1962). In poultry the hematologic response to corticosteroids is leukocytosis with heterophilia and lymphopenia (Hublé 1955, Glick 1961, Bell and Freeman 1971). However, it must be emphasized that the hematologic response to stress varies from species to species. For example, Bhat-tacharyya and Sarkar (1968) found that in the rock dove, the house crow (*Corvus splendens*) and the cattle egret (*Bubulcus ibis*) the response to cortical stimulation by ACTH and unilateral adrenalectomy was heteropenia and lymphocytosis. Only the common myna (*Acridotheres tristis*) responded in a similar fashion to poultry with heterophilia and lymphopenia in that study.

Levels of epinephrine and norepinephrine have been shown to be elevated during migration of the common snipe (*Gallinago gallinago*) and the rose-colored starling (*Sturnus roseus*; Epple and Stetson 1980). Information about the effect of catecholamines on avian hematology is lacking, but these substances are known to cause leukocytosis with neutrophilia and lymphocytosis in mammals (Duncan and Prasse 1986). Without having measured serum levels of catecholamines and corticosteroids during this study, it is impossible to determine the relative

level of stress for each bird. Birds with a total white blood cell count above the third quartile (leukocytic) had an absolute heterophilia, lymphocytosis and eosinophilia which may reflect more of a catecholamine-induced stress pattern rather than a corticosteroid-induced hemogram (which should present with lymphopenia). However, we caution against over-interpretation of the data since hormone levels were not measured and the effects of catecholamines and glucocorticoids on the hemogram are not known. Leukocytosis is often present with bacterial, fungal or parasitic infections, whereas leukopenia may accompany viral infections. Most hawks captured in this study appeared healthy on physical examination, showing no signs of clinical disease. Seriously ill birds might not be able to migrate and therefore would not be included in this study. Birds in this study with leukocytosis or leukopenia may have been ill, but the source of possible infection was not determined. We warn, however, that with lure traps a hungrier subset of the migrating population might have been sampled. This group may include birds who have no evidence of disease yet are subclinically ill and perhaps hungrier than healthy birds. But because no evidence of clinically significant disease states could be detected in the subjects, the white blood cell counts and other hematologic parameters for all birds from this study should be analyzed as a spectrum of hematologic findings for migrating, clinically healthy sharp-shinned hawks.

ACKNOWLEDGMENTS

The authors wish to thank the Wharton Trust, the Bailey Wildlife Foundation and the Hawk Migration Association of North America for providing support for this study. We thank Jerry Lehr and all the staff of the Little Gap Banding Station for assisting in this study. The Tufts Wildlife Clinic and its director, Charles J. Sedgwick, generously provided space and resources for laboratory and data analyses. Carolyn Haseltine lent her considerable expertise in avian hematology to the training of project personnel and interpretation of slides. This is Hawk Mountain Contribution number 21.

LITERATURE CITED

- ASHFORD, R. W., I. WYLLIE AND I. NEWTON. 1990. *Leukocytozoon toddi* in British sparrowhawks *Accipiter nisus*: observations on the dynamics of infection. *J. Nat. Hist.* 24:1101-1107.
- BALASCH, J., S. MUSQUERA, L. PALACIOS, M. JIMENEZ AND J. PALOMEQUE. 1976. Comparative hematology of some falconiforms. *Condor* 78:258-259.
- BELL, D. J. AND B. M. FREEMAN. 1971. Physiology and

bioch
New
BHATTA
leuk
hibit
BOND,
stud
non
CARPEN
altit
50A
CHRIST
Hän
ifvög
gung
Arte
COOPER
East
DAVIDA
asite
surv
DEIN, I
tolog
ton,
DUNCAN
orac
U.S.
ELLIOT
limir
Anin
EPPLE,
nolou
FERRER
J. C.
istry
migra
1123
GEE, G. I
Speci
crane
463--
GESSAM,
1986.
Coop
GLICK, P
desox
the v
Poult.
HALLIWF
WEDI

total white (leukocytic) osis and eoa catecholn a cortico-uld present ion against none levels cholamines not known. rial, fungal ia may accaptured in l examinae. Seriously id therefore irds in this y have been was not de- lture traps ation might clude birds ubclinically birds. But ant disease , the white parameters nalyzed as migrating,

Trust, the igration As- port for this of the Little . The Tufts lgwick, gen- oratory and onsiderable g of project is is Hawk

1990. *Leu-Accipiter ni-* tion. *J. Nat.*

A. JIMENEZ hematology

siology and

- biochemistry of the domestic fowl. Academic Press, New York, NY U.S.A.
- BHATTACHARYA, T.K. AND A.K. SARKAR. 1968. Avian leukocytic responses induced by stress & corticoid inhibitors. *Indian J. Exp. Biol.* 6:26-28.
- BOND, C.F. AND P.W. GILBERT. 1958. Comparative study of blood volume in representative aquatic and nonaquatic birds. *Am. J. Physiol.* 194:519-521.
- CARPENTER, F.L. 1975. Bird hematocrits: effects of high altitude and strength of flight. *Comp. Biochem. Physiol.* 50A:415-417.
- CHRISTOPH, H.J. AND G. BOROWSKI. 1961. Beiträge zur Hämatologie der Zootiere. IV. Das Blutbild von Greifvögeln (Accipitres) unter besonderer Berücksichtigung einger in Deutschland noch heimischer kleinerer Arten. *Kleintier-Praxis* 6:71-76.
- COOPER, J.E. 1975. Haematological investigations in East African birds of prey. *J. Wildl. Dis.* 11:389-394.
- DAVIDAR, P. AND E.S. MORTON. 1993. Living with parasites: prevalence of a blood parasite and its effect on survivorship in the purple martin. *Auk.* 110:109-116.
- DEIN, F.J. 1984. Laboratory manual of avian hematology. Association of Avian Veterinarians, Boca Raton, FL U.S.A.
- DUNCAN, J.R. AND K.W. PRASSE. 1986. Veterinary laboratory medicine. Iowa State Univ. Press, Ames, IA U.S.A.
- ELLIOTT, R.H., E.E. SMITH AND M. BUSH. 1974. Preliminary report on hematology of birds of prey. *J. Zoo Anim. Med.* 5:11-16.
- EPPLE, A. AND M.H. STETSON. 1980. Avian endocrinology. Academic Press, New York, NY U.S.A.
- FERRER, M., T. GARCÍA-RODRÍGUEZ, J.C. CARRILLO AND J. CASTROVIEJO. 1987. Hematocrit and blood chemistry in captive raptors (*Gyps fulvus*, *Buteo buteo*, *Milvus migrans*, *Aquila heliaca*). *Comp. Biochem. Physiol.* 87A: 1123-1127.
- GEE, G.F., J.W. CARPENTER AND G.L. HENSLER. 1981. Species differences in hematological values of captive cranes, geese, raptors, and quail. *J. Wildl. Manage.* 45: 463-483.
- GESSAMAN, J.A., J.A. JOHNSON AND S.W. HOFFMAN. 1986. Hematocrits and erythrocyte numbers for Cooper's and sharp-shinned hawks. *Condor* 88:95-96.
- GLICK, B. 1961. The effect of bovine growth hormone, desoxycorticosterone acetate, and cortisone acetate on the white blood cell counts of 2-week-old chickens. *Poult. Sci.* 40:1537-1539.
- HALLIWELL, W.H., G. IVINS, D.A. SCHMIDT AND G. WEDDLE. 1975. A preliminary report on the hematology and chemical profiles in selected birds of prey. *Proc. Ann. Meet. Am. Assoc. Zoo Vet.* 188-196.
- HUBLÉ, J. 1955. Haematological changes in cockerels after ACTH and cortisone-acetate treatment. *Poult. Sci.* 34:1357-1359.
- HUNTER, S.R. AND L.R. POWERS. 1980. Raptor hematocrit values. *Condor* 82:226-227.
- KIRKPATRICK, C.E. AND H.B. SUTHERS. 1988. Epizootiology of blood parasite infections in passerine birds from central New Jersey. *Can. J. Zool.* 66:2374-2382.
- LUMEIJ, J.T. AND J.J. DE BRUIJNE. 1985. Evaluation of the refractometric method for the determination of total protein in avian plasma or serum. *Avian Path.* 14: 441-444.
- OLSEN, G.H. AND S.D. GAUNT. 1985. Effect of hemoprotozoal infections on rehabilitation of wild raptors. *J. Am. Vet. Med. Assoc.* 187:1204-1205.
- PEIRCE, M.A. 1980. Current knowledge of the haematozoa of raptors. Pages 15-19 in J.E. Cooper and A.G. Greenwood [EDS.], Recent advances in the study of raptor diseases. Chiron Publications Ltd., West Yorkshire, U.K.
- PERRY, M.C., H.H. OBRECHT, B.K. WILLIAMS AND W.J. KUENZEL. 1986. Blood chemistry and hematocrit of captive and wild canvasbacks. *J. Wildl. Manage.* 50: 435-441.
- REHDER, N.B., D.M. BIRD, P.C. LAGUË AND C. MACKAY. 1982. Variation in selected hematological parameters of captive red-tailed hawks. *J. Wildl. Dis.* 18:105-109.
- SAS INSTITUTE, INC. 1988. SAS/STAT user's guide: Statistics. SAS Inst., Inc., Cary, NC U.S.A.
- SMITH, E.E. AND M. BUSH. 1978. Haematologic parameters on various species of Strigiformes and Falconiformes. *J. Wildl. Dis.* 14:447-450.
- SNYDER, J.E., D.M. BIRD AND P.C. LAGUË. 1980. Variations in selected parameters in the blood of captive American kestrels. Pages 113-115 in J.E. Cooper and A.G. Greenwood [EDS.], Recent advances in the study of raptor diseases. Chiron Publications Ltd., West Yorkshire, U.K.
- STURKIE, P.D. 1986. Body fluids: blood. Page 112 in P.D. Sturkie [ED.], Avian physiology. Springer-Verlag, New York, NY, U.S.A.
- WOLFORD, J.H. AND R.K. RINGER. 1962. Adrenal weight, adrenal ascorbic acid, adrenal cholesterol and differential leucocyte counts as physiological indicators of "stressor" agents in laying hens. *Poult. Sci.* 41:1521-1529.

Received 3 May 1993; accepted 7 May 1994