the synthesis of elvirol by the Australian workers on similar lines.

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## CHROMOSOME NUMBER AND DNA, RNA VALUES IN SOME INDIAN BATS (CHIROPTERA)

Our knowledge about the chromosomes of bats has been very meagre till todate despite their wide distribution. A survey of the literature reveals that cytogenetic reports of only a few Indian species of bats are available. The present studies were therefore, undertaken with a view to exploring the chromosomal data on as many representatives of bats as possible. The present findings, which are a part of a larger programme of research work on Indian bats, have been number of four species, viz., Rhinopoma microphyllum the most variable cytogenetic character while the perforatus perforatus and Hipposideros fulvus pallisidus larly, in the two species, viz., Hipposideros fulvus is being reported for the first time, whereas that for pallisidus (Microchireptera) and Cynopterus sphinx the remaining five species, viz., Cynopterus sphinx sphinx (Megachiroptera) both with a diploid number sphinx<sup>1</sup>, Rhinopoma hardwickei hardwickei<sup>1-2</sup>, Scoto- of 34 chromosomes, the DNA values are higher in philus heathi heathi<sup>3</sup>, Megaderma lyra lyra<sup>4</sup> and the former than in the latter. Similarly, Rhinopoma

also. The studies also include the estimation of DNA and RNA in mg/gm of known weight of tissues like liver and spleet in order to find out the possible relationship between these biochemical components of the nucleus and the chromosome number in the various species of bats under report.

For chromosomal studies, 0.5% of colchicine per kg body weight6 was injected intraperitoneally and 21 hours later the specimens were sacrificed and the marrow from the long bones was collected. After hypotonic treatment in sodium citrate (0.9%) for half an Four and fixation in acetic-alcohol (1:3) overnight6, the usual air-drying technique7 was followed.

The quantitative estimation of DNA and RNA contents from the tissues was confined to a known weight (100 mg) by employing perchloric acid method<sup>8-9</sup>. The readings for DNA and RNA were taken from at least five different samples of the same species for every tissue on Bausch and Lomb Spectronic-20. The mean values in mg/gm were thus calculated from the standard graphs prepared by using the standard DNA's and RNA's of calf thymus gland and yeast respectively. The standard deviations and standard errors of the 'mean' were also calculated which refer to variation among the averages obtained from different individuals of the same species.

The studies of chromosomal slides reveal the diploid number of chromosomes varying from 34-54 in these various species (Table I). It is 34 in two, 42 in two. 36 in four and 54 in one species. The studies point cut that the 2N number of chromosomes varies not only in the species belonging to different genera but also in the species of the same genus. Thus the species are rather indistinguishable on the basis of chremosome number alone.

The mean values for DNA, RNA in mg/gm. of the known weight of the various ti-sues obtained through repeated experimentation are also mentioned in Table I and Figs. 1 and 2.

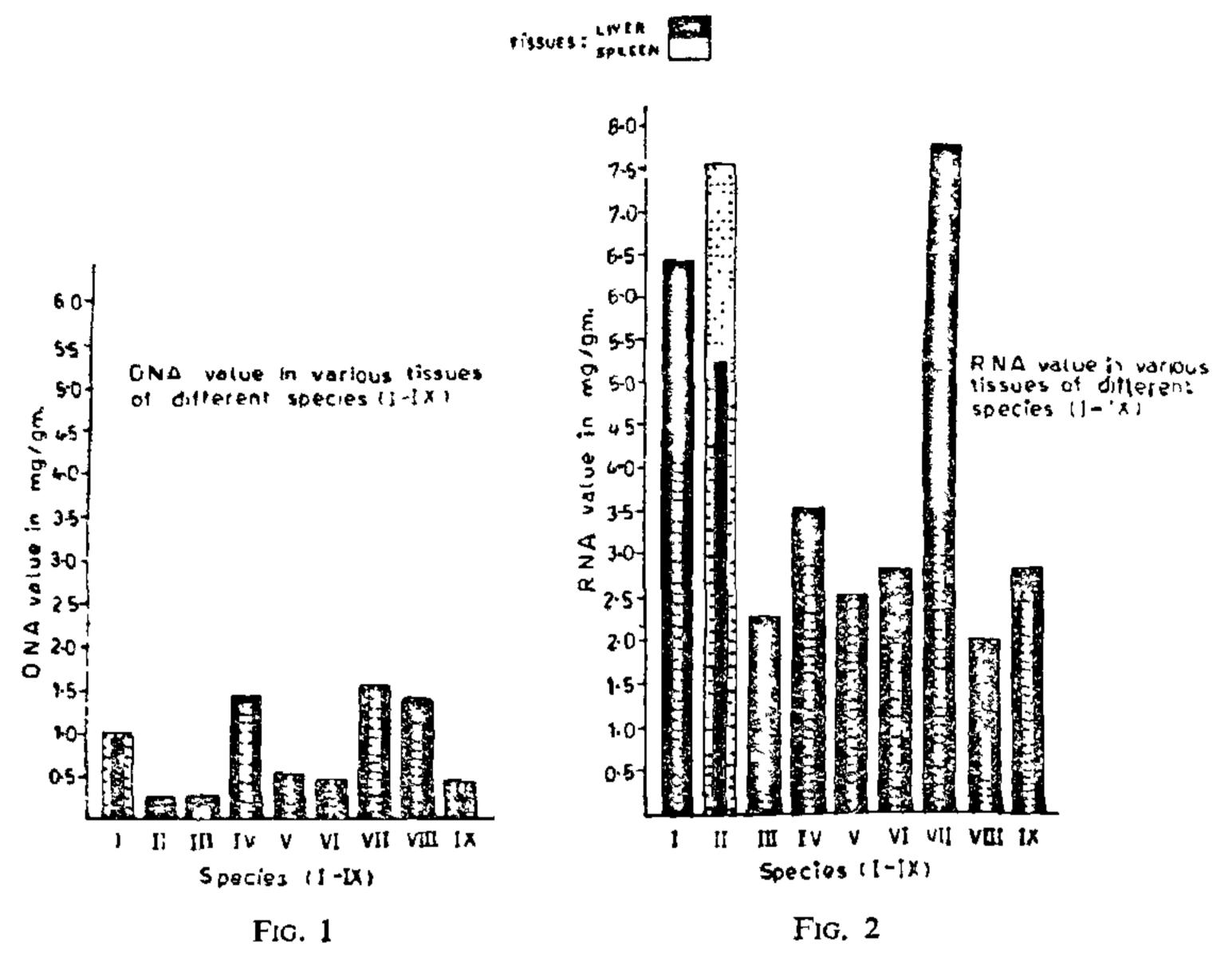
Considering the chromosome number and DNA and RNA values on a collective basis, one finds that the species with the same diploid number of chromosomes possess a variable amount of DNA and RNA in their tissues. This has also been reported in the various species of the genus Bufo and in Amphibia made on rine species. Of these, the chromosomal in general10 where the total nuclear DNA amount is kinneari, Taphozous nudiventris kachensis, Taphozous chromosome numbers are relatively constant. Simi-Rousettus leschenaulti5 had been known beforehand hardwickei hardwickei, Taphozous nudiventris kachensis

TABLE I

nucleic acid (DNA) and ribose nucleic acid (RNA) values in mg/gm with standard deviation (S.D.) standard error (S.E.) in nine species\* of bats and deoxyribose Name, locality, diploid number

			314,	standard error (3.12.) in nine species	ecies of pars		
Sr.	Names of the Species	Locality	Diploid	DNA value (mg/gm) ± S.D.± S.E	n) ± S.D. ± S.E.	RNA value (mg/g	lue (mg/gm) ± S.D. ± S.E.
				Liver	Spleen	Liver	Spleen
Order C Suborder Family—I Cynopteru	Order Chiroptera Suborder Megachiroptera Family—Pteropodidae Cynopterus sphinx sphinx						
(Vahl) II. Rousettu:	(Vahl) Rousettus leschenaulti	Ambala	34	0.960±0.054±0.027	1.005±0.017±0.008	6.400±0.200±0.090	4.200±0.240±0.110
	CDesmarest) Suborder Microchiroptera Family—Rhinopomatidae Rhinopoma microphyllum	do.	36	0.240±0.040±0.020	0.180±0.020±0.010	5.200±0.200±0.090	7.500±0.230±0.110
kinn IV. Rhinop	kinneari Wroughton Rhinopoma hardwickei	Agra	42	0·270±0·050±0·020	0.105±0.005±0.002	2·250±0·440±0·200	0.225±0.090±0.040
hard Family V. Tapho:	hardwickei Family—Emballonuridae Taphozous nudiventris	Jodhpur	36	1-395±0-0005±0-0004	1.380±0.081±0.030	3.500±0.014±0.007	3.000±0.500±0.230
kach VI. Taphoz	kachensis Taphozous perforatus	Ambala	36	0.510±0.060±0.030	0.536±0.001±0.0005	2.470±0.750±0.340	2·500±0·100±0·040
perfo Family VII. Megad	perforatus Family—Mcgadermatidae Megaderma lyra lyra	Jodhpur	45	0.490±0.230±0.100	0.460±0.160±0.070	2⋅800±0⋅150±0⋅070	2.050±0.050±0.020
(Geo Family WIII. Hippos	(Geoffroy)  Family—Hipposideridae  Hipposideros fulvus	Ambala and Agra	24	1.530±0.150±0.070	1.530土0.040土0.010	7.700±0.230±0.100	3.400±0.100±0.040
pallisi Family-	pallisidus amily—Vespertilionidae	Jodppur	34	1.360±0.150±0.070	1.270±0.100±0.040	2.000±0.100±0.040	1.000±0.200±0.090
IX. Scotop	7	Ambala	36	0.250±0.040±0.020	0.380±0.010±0.005	2.800±0.270±0.100	2·600±0·350±0·160

\* The species have been identified by Zoological Survey of India, Calcutta.



Figs. 1-2. DNA, RNA values in different species (I-IX) of bats.

Species: I—Cynopterus sphinx sphinx; II—Rousettus leschenaulti; III—Rhinopoma microphyllum kinneari; IV—Rhinopoma hardwickei hardwickei; V—Taphozous nudiventris kachensis; VI—Taphozous perforatus perforatus; VII—Megaderma lyra lyra; VIII—Hipposideros fulvus pallisidus; IX—Scotophilus heathiheathi.

and Scotophilus heathi heathi each with 36 chromosomes in their diploid sets possess different amount of DNA and RNA. This is also true for Rhinopoma microphyllum kinneari and Taphozous perforatus perforatus having diploid number of 42 chromosomes. The studies thus suggest that the amount of DNA and RNA in a tissue of a species forms a distinctive and a characteristic parameter for differentiating the species having the same number of chromosomes in their diploid sets.

The present findings also reveal that except for Rousettus leschenaulti, Rhinopora micrephyllim and Scotophilus heathi heathi, the DNA values in the rest of the species are nearly constant in the two tissues dealt with. This, however, does not hold true for the RNA values which vary not only in the two tissues of the same species but also in a specific tissue of different species.

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## ISOLATION OF CORYNEBACTFRIUM PYOGENES FROM AN ABORTED EQUINE FOETUS

The bacterium Corynebacterium pyogenes, first described by Lucet in 1893 from suppurative lesions in cattle, is widespread amongst domestic animals affecting cattle, sheep, goats and pigs. The common disease conditions caused by this organism are suppurative pneumonia, suppurative arthritis and other