



Observation on Proteinuria in Cases of Kala-Azar (Visceral Leishmaniasis)

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ABSTRACT

Kala-azar has been endemic in the eastern part of India since more than a century. The principal area affected are North Bihar, West Bengal, Assam, Tripura, Eastern U.P. & Nepal. In this study cases admitted in P.M.C.H were studied for proteinuria it was observed that 44% kala-azar cases had proteinuria and 18% proteinuria with haematuria.

INTRODUCTION

Visceral leishmaniasis is a tropical disease produced by protozoan *Leishmania donovani* characterised by irregular fever, splenohepatomegaly, weight loss, anemia, leucopenia & hyperglobinaemia, endemic in Bihar, West Bengal, & Eastern part of India.

In patient with kala-azar, occasionally renal lesions has been described but with little disturbance of renal function. Slight albuminuria has been described regularly in kala-azar patients (Cole ACE 1944 in East Africa, Von Peenen & Reid 1962 from Upper Wile Province, Maru 1979 in North Western Ethiopia) In Sudan (Sati 1962) albuminuria was reported in 50% of Brazilian patients elevated urea & creatinine are only seen terminally. Microscopic haematuria is also frequent.

Although several foreign workers have shown abnormal urine analysis in visceral leishmaniasis still not much work has been done in case of Indian kala-azar.

MATERIALS & METHODS

A total number of 100 patients of Kala-azar admitted to Kala-azar ward & medical indoor ward of Patna Medical College & Hospital were subject to present study. Diagnosis of Kala-azar was confirmed clinically as well as demonstrating parasite in bone marrow or splenic puncture smear. A thorough clinical examination of each patient was done and any another disease as Diabetis Mellitus, Jaundice, Cirrhosis of liver, Pulmonary tuberculosis, Pneumonia, Cardiac disease, Hypertension were excluded from the study. Only the fresh cases of Kala-azar without previous history of renal pathology were included in our study.

All confirmed cases of Kala-azar were subjected to following renal function test:-

Routine examination of Urine especially for albumin, cells and casts.

Estimation of Blood Urea, Serum Creatinine.

Estimation of GFR (by endogenous creatinine clearance).

24 hour total Urinary protein excretion.
USG of whole abdomen.
Other investigation as TC-DC of WBC, ESR, X-Ray chest, LFT, Viral markers to exclude other pathology.

OBSERVATION

Hundred cases of kalazar patient were studied for renal function.

Table 1 Showing Age wise distribution.

S.N	AGE	No of cases	Percentage	Mean (year)
1.	11 - 20	54	54	16.04
2.	21 - 30	36	36	26.55
3.	31 - 40	04	04	35.00
4.	41 - 50	06	06	43.66

Table 2 Showing Sex wise distribution

S.N	Sex	No of cases	Percentage	Male : Female ratio
1.	Male	70	70	7 : 3
2.	Female	30	30	

Table 3 Showing duration of illness in Patients

Duration illness (Weeks)	No of Cases	Mean weeks	Percentage
1 – 8	34	7.37	34
9 – 16	48	13.31	48
17 – 29	18	21.33	18

Table 4 Showing urine analysis

Urinary findings	No of Cases	Percentage
Proteinuria	22	44
Haematuria + Proteinuria	09	18

Table 5 Showing quantitative Proteinuria

Total Urine Protein in 24hr range (mg/dl)	No of Patients	Mean urine protein in 24 hr range
Upto -50	32	22.35
50 – 100	12	95.35
100 – 300	42	298.28
300 – 1 Gram	11	695.68
Above 1 Gram	03	1193.66

DISCUSSION

The present work observation on proteinuria in cases of kala-azar was carried out in patients admitted in Nephrology Unit and kala-azar ward. On of 100 cases 70 were male and 30 were females and mean age was 22.24.

In the present study no patients has the nephritic syndrome. Abnormal urinalysis was observed in

52% of cases, out of which 44% has proteinuria and 18% has proteinuria with haematuria (In sudan sati 1962 albuminuria was reported in 50% cases) The incidence of proteinuria in the present series was 44% in Indian kala-azar patients thought majority of patients did not develop proteinuria. Proteinuria might be renal pathology due to urinary tract lesions or renal pelvis, bladder, prostate but all the lesion were excluded before assessing renal states, however fever and anemia which were main presentation of kala-azar might have secondary effects on kidney producing proteinuria but in this conditions significant proteinuria did not occur. It was difficult to opine that proteinuria present in some patients of kala-azar was an integral part of disease or not. Duration of illness seemed to have some association with proteinuria.

CONCLUSION

In the present study it has been documented that proteinuria has been observed frequently in kala-azar patients. Further study are needed to document the specificity of the glomerular, tubular & intestinal lesions observed in parasitic diseases.

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