

# URINARY MONOCYTE CHEMOTACTIC PROTEIN-1 IS ASSOCIATED WITH URINARY OXIDATIVE STRESS, MICROALBUMINURIA AND THE BACTERIOLOGICAL INDEX IN LEPROSY

Geraldo Bezerra da Silva Júnior <sup>2,3</sup>; Gdayllon Cavalcante Meneses <sup>1</sup>; Marcus Felipe Bezerra da Costa <sup>1</sup>; Heitor de Sá Gonçalves <sup>4</sup>; Elizabeth De Francesco Daher <sup>2</sup>; Alexandre Braga Libório <sup>2</sup>; Alice Maria Costa Martins <sup>1</sup>.

<sup>1</sup>Post-Graduation Program in Pharmacology, Federal University of Ceará. Fortaleza, Ceará, Brazil. <sup>2</sup>Department of Internal Medicine, School of Medicine, Federal University of Ceará. Fortaleza, Ceará, Brazil. <sup>3</sup>School of Medicine, Health Sciences Center, University of Fortaleza. Fortaleza, Ceará, Brazil. <sup>4</sup>Division of Dermatology, Centro de Dermatologia Dona Libania. Fortaleza, Ceará, Brazil.

## OBJECTIVES

Renal lesions in leprosy have been extensively described in medical literature. Leprosy patients can present with kidney disease from glomerular (glomerulonephritis, amyloidosis) or tubule-interstitial etiology. The aim of this study is to evaluate renal abnormalities in leprosy patients through traditional biomarkers of renal disease and Monocyte Chemotactic Protein-1 (MCP-1).

## METHODS

This is a cross-sectional study of 44 patients with clinical and laboratory diagnosis of leprosy with no previous anti-mycobacterium treatment. Patients were recruited in public health centers in Fortaleza, Ceara, Brazil, between August 2012 and August 2013. A group of 15 healthy subjects were included as a control group. Skin smear was assessed through a bacteriological index - from 0 to 6+. Glomerular filtration rate (GFR), protein excretion, microalbuminuria, urinary oxidative stress (malondialdehyde-MDA) and urinary MCP-1 were estimated. All urine measurements were normalized by urinary creatinine concentration.

## RESULTS

Table 1: Comparison of renal function parameters between the clinical forms of leprosy and control groups.\*p<0.05(DV/VV vs TT/DT and Control).\*\*p<0.05(TT/DT, DD, DV/VV vs Control)

Parâmetros	TT/DT (n=14)	DD (n=19)	DV/VV (n=11)	Control (n=15)
S <sub>Cr</sub> (mg/dL)	0,85±0,21	0,89±0,12	0,80±0,17	0,81±0,14
GFR <sub>E</sub> mL/min/1.73m <sup>2</sup>	124±31	107±23	117±35	115±18
Urine Protein Excretion (mg/g-Cr)	60±29	86±52	<b>129±72*</b>	62±37
Albumin Excretion (mg/g-Cr)	4,4±3,9	4,5±3,7	5,1±3,7	2,0±1,2
S <sub>Na</sub> (mEq/L)	142±2,8	141±2,6	139±3,8	139±1,5
S <sub>K</sub> (mEq/L)	4,5±0,3	4,4±0,4	4,5±0,7	4,4±0,4
FE <sub>Na</sub> (%)	0,7±0,5	0,8±0,6	0,7±0,5	0,5±0,3
FE <sub>K</sub> (%)	4,4±2,3	6,9±5,1	6,8±5,4	5,6±3,0
PCR	1,4±0,9	6,0±13	22,9±33,4	---
MDA (mmol/g-Cr)	<b>1,7±1,4**</b>	<b>1,5±0,7**</b>	<b>2,3±1,7**</b>	0,12±0,07

Figure 2. Pearson correlations of urinary (MCP-1) with urinary MDA and urine albumin excretion. \* p<0.05, \*\*p<0.01.

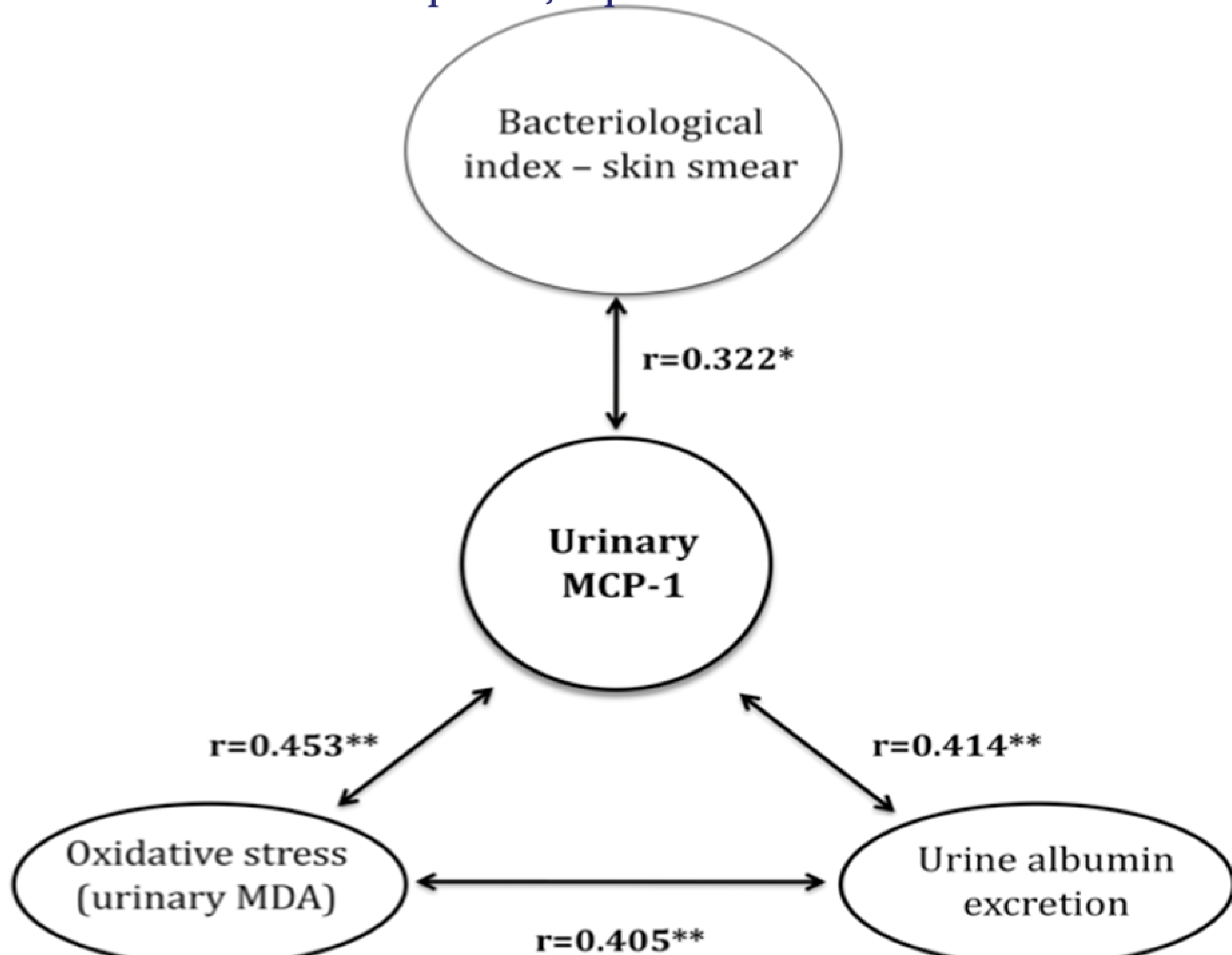
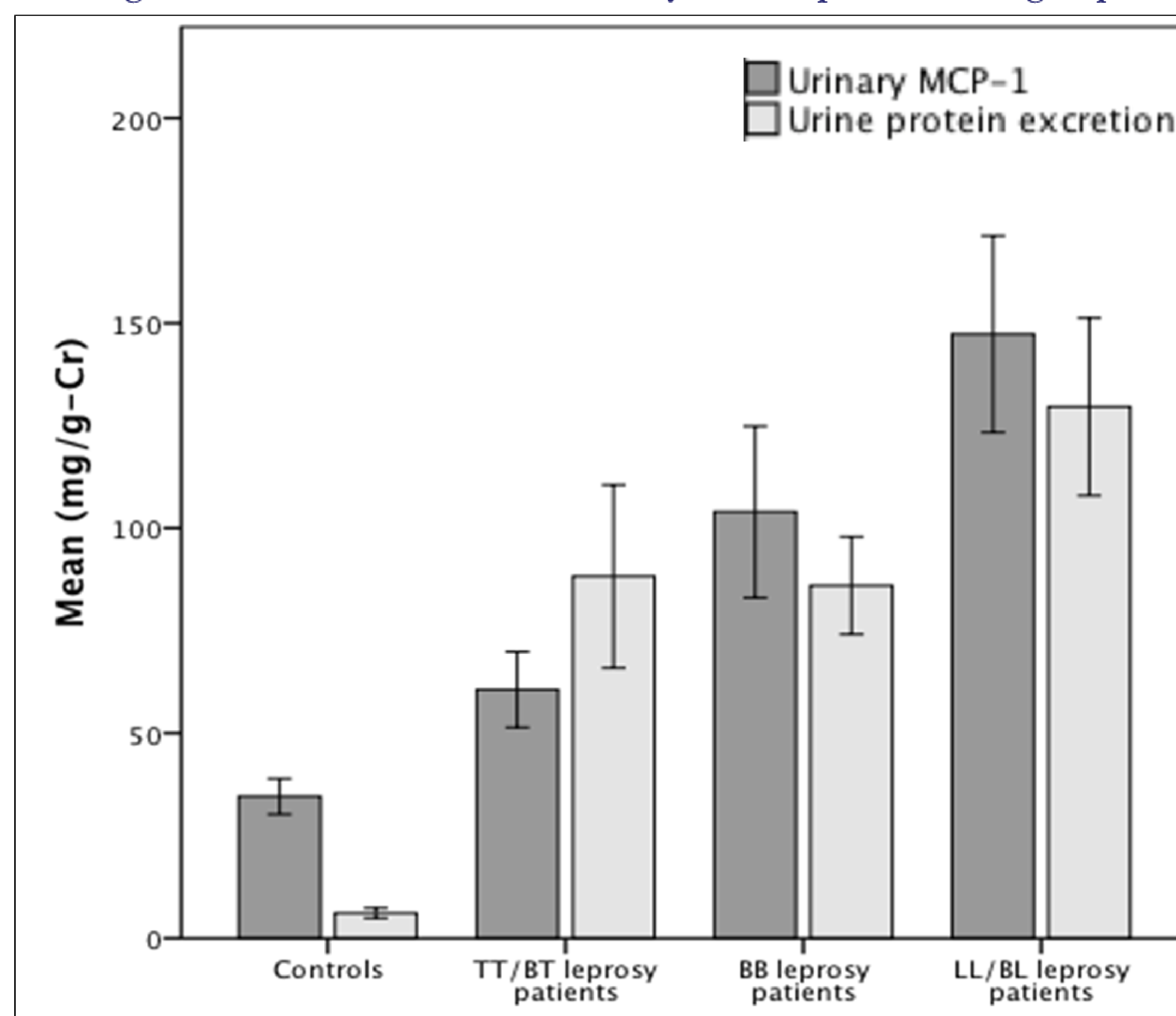


Figure 1: Urinary MCP-1 and protein excretion in controls and leprosy patients according their clinical classification. Urinary MCP-1: p<0.05 for all groups.



## CONCLUSION

We demonstrated that leprosy patients with no clinical kidney disease have increased urinary MCP-1 and its levels are even higher as patients approximates to lepromatous polar form. Moreover, urinary MCP-1 was associated with urinary oxidative stress and urine albumin excretion, suggesting that these patients are at increased risk of developing clinical kidney disease in the future.

## REFERENCES

- Kim MJ, Tam FW. Urinary monocyte chemoattractant protein-1 in renal disease. *Clin Chim Acta* 2011; 412: 2022-2030.
- Oliveira RA, Silva GB Jr, Souza CJ et al. Evaluation of renal function in leprosy: a study of 59 consecutive patients. *Nephrol Dial Transplant* 2008; 23: 256-262.
- Daher EF, Silva GB Jr, Cezar LC et al. Renal dysfunction in leprosy: a historical cohort of 923 patients in Brazil. *Trop Doct* 2011; 41: 148-150.

E-mail: geraldobezerrajr@yahoo.com.br, gdayllon@yahoo.com.br, efdaher@uol.com.br.