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Metabolic Profile and Cardiovascular Risk in Patients with Glomerular diseases treated with Immunosuppressants

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OBJECTIVES

Immunosuppressants (IS) are often required

to treat glomerulonephritis (GN) to prevent

This was a retrospective cohort study of adults with biopsy-proven GN diagnosed between

METHODS

13th January 2011 and 28th July 2015. Patients <21 years and those who received IS prior to

subsequent renal failure. However,
metabolic complications arising from IS
therapy may also lead to undesirable
outcomes such as cardiovascular disease
(CVD). We compared diabetics (DM) and
non-diabetics who received IS therapy for
GN and evaluated risk factors for CVD.

kidney biopsy were excluded.

Demographic, comorbidity, clinical and pharmacotherapy data were retrieved from electronic medical records. Pre-biopsy fasting glucose, triglyceride (TG), low-density lipoprotein (LDL) and high-density lipoprotein (HDL) within 6 months preceding kidney biopsy and peak values within 6 months after IS therapy were obtained.

CVD was present if there was an admission for acute myocardial infarction, congestive cardiac failure, or if coronary angiogram found >50% stenosis in the coronary vessels.

	Diabetic N = 41	Non-diabetic N = 244	P value
Age at diagnosis, years	55.9 (45.9 <i>,</i> 66.3)	47.1 (33.6, 62.1)	0.003
Male, n (%)	23 (56.1)	104 942.6)	0.11
Comorbidities, metabolic prof	file and medications befo	ore biopsy	
Hypertension, n (%)	32 (78.0)	100 (41.0)	<0.001
Dyslipidemia, n (%)	26 (63.4)	57 (23.4)	<0.001
Serum creatinine, µmol/L	124 (84, 197)	107 (73 <i>,</i> 189)	0.19
UPCR, g/g	5.38 (3.03, 11.51)	5.45 (2.41, 9.35)	0.23
aGlucose	6.2 (5.4, 8.1)	5.2 (4.8, 5.8)	<0.001
^a TG, mmol/L	1.61 (1.21, 2.66)	1.81 (1.21, 2.36)	0.71
^a LDL, mmol/L	3.29 (4.14, 2.26)	4.04 (2.80, 6.52)	0.02
^a HDL, mmol/L	1.07 (0.88, 1.37)	1.34 (1.10, 1.77)	0.006
Anti-lipid medications	33 (80.5)	94 (38.5)	<0.001
Clinical and laboratory parameters after immunosuppressive therapy			
Change in ^a glucose, %	46.9 (3.2, 80.8)	12.4 (0, 38.6)	0.003
Change in ^a TG, %	18.6 (-19.9, 53.2)	-8.9 (-40.9, 25.5)	0.03
Change in ^a LDL, %	-3.8 (-40.2, 52.2)	-15.1 (-55.6, 16.5)	0.16
Change in ^a HDL, %	15.6 (-7.2, 35.2)	18.8 (-14.1, 46.0)	0.73

^aAll glucose and lipid values were obtained from fasting samples

UPCR, urine protein to creatinine ratio; TG, triglyceride; LDL, low density lipoprotein; HDL, high density lipoprotein; CVD, cardiovascular disease

Categorical variables are expressed as number (percentage) and compared using chi-square or Fisher's exact test as appropriate. Continuous variables are expressed as median (interquartile range) and compared using Mann-Whitney U test.

(26.7%), IgA nephropathy (18.6%), membranous nephropathy (14.4%) ses. 3.2, 51.4) months. of diabetics and non-diabetic patients Patients with DM were older, more idemia than non-diabetics. Diabetics doses [30 (20, 50) mg vs. 50 (30, 60) eceived cyclosporine (36.6% vs. 12.3%, tients with DM had greater increase in were morely likely to have CVD, than non-diabetics. There was no difference in follow up duration [24.6] (13.0, 56.9) months vs. 29.4 (13.1, 50.7) months, p>0.05].



CONCLUSION

CVD occurred in 14 patients at 11.7 (4.2, 28.2) months after biopsy.

After adjusting for age, gender, comorbidities, renal function, proteinuria and post-IS therapy levels of glucose and lipid, DM [adjusted OR 4.98 (95% CI: 1.18, 21.01), p=0.03] was independently associated with CVD.

Diabetics with GN were more likely than non-

diabetics to have concomitant cardiovascular risk

factors and increased glucose and lipid levels

after IS therapy. DM was an independent risk

factor for CVD after IS therapy.

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