Recurrence Of Focal Segmental Glomerulosclerosis Among Kidney Transplant Recipients: Kuwait Experience

Authors:

Osama Gheith, Zakaryia Elsayed, Zoheer Fayyad, Mohamed Shaker, Medhat Alawady, Nabil Alserwy, Prasad Nair, Khaled Abdultawab, Mohamed Balaha, Ayman Nagib, Hasaneen Aboatteya, and Torki AlOtaibi.





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Background and aim of the study

• Post-transplant FSGS is a major risk factor for graft loss. We therefore aimed to assess patients with FSGS in our cohort of kidney transplant recipients.

Patients and methods

- In this retrospective study that aimed to investigate glomerular disease recurrence post-transplantation.
- Kidney transplant recipients (KTR) were screened for the diagnosis of FSGS between 1996 and 2023 and details were recorded about the transplant, clinical outcomes, treatments, and other risk factors.

Results

- Among 3670 KTR screened for FSGS, 106 were identified to have FSGS as original kidney disease.
- Of them, 52 had a diagnosis of idiopathic FSGS. Most of the patients were Kuwaiti males who received their grafts from living donors (84.9%). Prophylactic plasma exchange (PE) was performed in 19 (38%). All patients were maintained on CNI based triple immunosuppression.

Results

- FSGS recurrence was confirmed in 10 patients (17.5%). PE and rituximab were the most frequent treatment options (7 out of 10 patients received PE and rituximab) and 4 patients had recurrence despite prophylactic PE and rituximab. The remaining patients were managed by anti-proteinuric agents.
- At one year follow-up, complete remission was observed in 5 patients while the remaining showed partial remission. Two patients lost their grafts within 2 years while the remaining 8 are enjoying stable graft function. Patient or graft outcomes were comparable between primary and secondary FSGS cases(P>0.05).

Table 1: Showed the some of the demographics of the studied population.

	1ry FSGS(N=57)	2 ^{ndry} FSGS (N=52)	P value
	Number %	Number %	
Mean age at transplant(years)	40.9±15.3	39.7±18.9	0.04
Dialysis mode			
Preemptive	14(24.6)	12(23.1)	
Hemodialysis	38(66.8)	37(71.1)	
Peritoneal dialysis	5(8.8)	3(5.8)	0.42
Immunosuppression			
Induction:			
None	8(14)	6(11.5)	
Basilixmab	17(29.8)	9(17.3)	
Lymphocyte depleting agents	23(40.4)	22(42.3)	
Others	9(15.8)	15(28.8)	0.38
Immediate graft function			
Immediate	40(72.7)	28(54.9)	
Slow	9(16.4)	12(23.5)	
DGF	4(7.3)	4(7.8)	
Unknown	2(3.6)	7(13.7)	0.159

Table 2: Showed post-transplant complications in the studied patients

Variables:	1ry FSGS(N=57)	2 nd FSGS (N=52)	P value
variables:	Number %	Number %	r value
Post-transplant complications:			
BK viremia	9(17.3)	7(14.0)	0.646
BK nephropathy	3(5.7)	1(2.0)	0.337
CMV viremia	10(23.8)	15(40.5)	0.111
NODAT	14(26.4)	12(24)	0.77
Mean rejection episodes	1.71±1.68	1±1	0.22
Graft outcome	50(87.7)	43(82.7)	
Functioning	7(12.3)	9(17.3)	0.45
Failed	7(12.3))(17.5)	U.TJ
Patient outcome			
Living	57(100)	48(92.3)	
Dead	0(0)	2(3.8)	
Lost follow up	0(0)	2(3.8)	0.11

Table 3: Showed numerical parameters of the studied patients

Variables	No recurrence FSGS (N=47)	Recurrent FSGS (N=10)	P value
	Number ± SD	Number ± SD	
FSGS treatment"	21(44.7)	4(40)	0.78
ACEi/ARB (number / %)		, ,	
PE (number / %)	17(36.2)	7(70)	0.049
Rituximab (number / %)	11(23.4)	7(70)	0.004
PE sessions(mean/SD)	6.3(2.8)	11(8.03)	
Pre-transplant weight (kg)			
Plasma exchanges	17(36.2)	7(70)	0.049
PE number	6.33±2.8	11±8.03	0.38
Weight at last follow up(kg)			

Table 4: Showed follow up parameters of the studied patients

Variables	No recurrence FSGS (N=47)	Recurrent FSGS (N=10)	P value
	Number ± SD	Number ± SD	
Serum creatinine (umol/L)			
Basal	124±77	150±73	0.33
3 months	108±38	117±35	0.48
6 months	106±39	189±160	0.003
1year	107±38	192±104	0.003
Last follow up	144±125	202±144	0.24
Proteinuria (g/24hours)			
Basal	0.91±0.82	1.3±1	0.205
3 months	0.26±0.44	1.5±2	<0.001
6 months	0.31±0.66	1±1.22	0.019
1year	0.37±0.71	1±1.5	0.059
Last follow up	0.79±1	1.9±1.1	0.004

Conclusions

•We reported lower rate of recurrence of idiopathic FSGS in our cohort (15.3 %) and the comparable patient and graft outcomes might be due to the our adopted optimized immunosuppressive regimen.