

World Journal of *Nephrology*

Quarterly Volume 13 Number 1 March 25, 2024



EDITORIAL

Raikou VD. Renoprotective strategies. *World J Nephrol* 2024; 13(1): 89637 [DOI: [10.5527/wjn.v13.i1.89637](https://doi.org/10.5527/wjn.v13.i1.89637)]

Sabath E. Point of care ultrasonography as the new “Laennec Sthetoscope”. *World J Nephrol* 2024; 13(1): 90542 [DOI: [10.5527/wjn.v13.i1.90542](https://doi.org/10.5527/wjn.v13.i1.90542)]

OPINION REVIEW

Peticca B, Prudencio TM, Robinson SG, Karhadkar SS. Challenges with non-descriptive compliance labeling of end-stage renal disease patients in accessibility for renal transplantation. *World J Nephrol* 2024; 13(1): 88967 [DOI: [10.5527/wjn.v13.i1.88967](https://doi.org/10.5527/wjn.v13.i1.88967)]

ORIGINAL ARTICLE

Retrospective Study

Jafry NH, Manan S, Rashid R, Mubarak M. Clinicopathological features and medium-term outcomes of histologic variants of primary focal segmental glomerulosclerosis in adults: A retrospective study. *World J Nephrol* 2024; 13(1): 88028 [DOI: [10.5527/wjn.v13.i1.88028](https://doi.org/10.5527/wjn.v13.i1.88028)]

Juarez-Villa JD, Zepeda-Quiroz I, Toledo-Ramírez S, Gomez-Johnson VH, Pérez-Allende F, Garibay-Vega BR, Rodríguez Castellanos FE, Moguel-González B, Garcia-Cruz E, Lopez-Gil S. Exploring kidney biopsy findings in congenital heart diseases: Insights beyond cyanotic nephropathy. *World J Nephrol* 2024; 13(1): 88972 [DOI: [10.5527/wjn.v13.i1.88972](https://doi.org/10.5527/wjn.v13.i1.88972)]

SYSTEMATIC REVIEWS

Ndongo M, Nehemie LM, Coundoul B, Diouara AAM, Seck SM. Prevalence and outcomes of polycystic kidney disease in African populations: A systematic review. *World J Nephrol* 2024; 13(1): 90402 [DOI: [10.5527/wjn.v13.i1.90402](https://doi.org/10.5527/wjn.v13.i1.90402)]

ABOUT COVER

Peer Reviewer of *World Journal of Nephrology*, Amgad E El-Agroudy, MBBCh, MD, FACP, FASN, FAST, Internal Medicine and Nephrology, College of Medicine and Medical Sciences, Arabian Gulf University, Manama, Bahrain. amgadelagroudy@hotmail.com

AIMS AND SCOPE

The primary aim of *World Journal of Nephrology* (WJN, *World J Nephrol*) is to provide scholars and readers from various fields of nephrology with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJN mainly publishes articles reporting research results obtained in the field of nephrology and covering a wide range of topics including acute kidney injury, acute or chronic interstitial nephritis, AIDS-associated nephropathy, anuria, chronic kidney disease and related complications, CKD-MBD, diabetes insipidus, diabetic nephropathies, Fanconi syndrome, glomerular diseases, inborn or acquired errors renal tubular transport, renal hypertension, kidney cortex necrosis, renal artery obstruction, renal nutcracker syndrome, renal tuberculosis, renal tubular acidosis, thrombotic microangiopathy, uremia, and Zellweger syndrome, *etc.*

INDEXING/ABSTRACTING

The WJN is now abstracted and indexed in PubMed, PubMed Central, Reference Citation Analysis, China Science and Technology Journal Database, and Superstar Journals Database.

RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: *Yu-Qing Zhao*; Production Department Director: *Xu Guo*; Editorial Office Director: *Ji-Hong Liu*.

NAME OF JOURNAL

World Journal of Nephrology

ISSN

ISSN 2220-6124 (online)

LAUNCH DATE

February 6, 2012

FREQUENCY

Quarterly

EDITORS-IN-CHIEF

Li Zuo, Ying-Yong Zhao

EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/2220-6124/editorialboard.htm>

PUBLICATION DATE

March 25, 2024

COPYRIGHT

© 2024 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/gerinfo/287>

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

PUBLICATION ETHICS

<https://www.wjgnet.com/bpg/gerinfo/288>

PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/gerinfo/239>

ONLINE SUBMISSION

<https://www.f6publishing.com>



Retrospective Study

Clinicopathological features and medium-term outcomes of histologic variants of primary focal segmental glomerulosclerosis in adults: A retrospective study

Nazarul Hassan Jafry, Shumaila Manan, Rahma Rashid, Muhammed Mubarak

Specialty type: Urology and nephrology

Provenance and peer review: Invited article; Externally peer reviewed.

Peer-review model: Single blind

Peer-review report's scientific quality classification

Grade A (Excellent): 0
Grade B (Very good): 0
Grade C (Good): C, C
Grade D (Fair): 0
Grade E (Poor): 0

P-Reviewer: Tanaka H, Japan

Received: September 6, 2023

Peer-review started: September 6, 2023

First decision: November 21, 2023

Revised: November 30, 2023

Accepted: January 11, 2024

Article in press: January 11, 2024

Published online: March 25, 2024



Nazarul Hassan Jafry, Shumaila Manan, Department of Nephrology, Sindh Institute of Urology and Transplantation, Sindh, Karachi 74200, Pakistan

Rahma Rashid, Muhammed Mubarak, Department of Pathology, Sindh Institute of Urology and Transplantation, Sindh, Karachi 74200, Pakistan

Corresponding author: Muhammed Mubarak, MD, Professor, Department of Pathology, Sindh Institute of Urology and Transplantation, Chand Bibi Road, DPMC, Sindh, Karachi 74200, Pakistan. drmubaraksiut@yahoo.com

Abstract

BACKGROUND

The Columbia classification identified five histological variants of focal segmental glomerulosclerosis (FSGS). The prognostic significance of these variants remains controversial.

AIM

To evaluate the relative frequency, clinicopathologic characteristics, and medium-term outcomes of FSGS variants at a single center in Pakistan.

METHODS

This retrospective study was conducted at the Department of Nephrology, Sindh Institute of Urology and Transplantation, Karachi, Pakistan on all consecutive adults (≥ 16 years) with biopsy-proven primary FSGS from January 1995 to December 2017. Studied subjects were treated with steroids as a first-line therapy. The response rates, doubling of serum creatinine, and kidney failure (KF) with replacement therapy were compared between histological variants using ANOVA or Kruskal Wallis, and Chi-square tests as appropriate. Data were analyzed by SPSS version 22.0. P -value ≤ 0.05 was considered significant.

RESULTS

A total of 401 patients were diagnosed with primary FSGS during the study period. Among these, 352 (87.7%) had a designated histological variant. The not otherwise specified (NOS) variant was the commonest, being found in 185 (53.9%) patients, followed by the tip variant in 100 (29.1%) patients. Collapsing (COL), cellular (CEL), and perihilar (PHI) variants were seen in 58 (16.9%), 6 (1.5%), and 3

(0.7%) patients, respectively. CEL and PHI variants were excluded from further analysis due to small patient numbers. The mean follow-up period was 36.5 ± 29.2 months. Regarding response rates of variants, patients with TIP lesions achieved remission more frequently (59.5%) than patients with NOS (41.8%) and COL (24.52%) variants ($P < 0.001$). The hazard ratio of complete response among patients with the COL variant was 0.163 [95% confidence interval (CI): 0.039-0.67] as compared to patients with NOS. The TIP variant showed a hazard ratio of 2.5 (95%CI: 1.61-3.89) for complete remission compared to the NOS variant. Overall, progressive KF was observed more frequently in patients with the COL variant, 43.4% ($P < 0.001$). Among these, 24.53% of patients required kidney replacement therapy ($P < 0.001$). The hazard ratio of doubling of serum creatinine among patients with the COL variant was 14.57 (95%CI: 1.87-113.49) as compared to patients with the TIP variant.

CONCLUSION

In conclusion, histological variants of FSGS are predictive of response to treatment with immunosuppressants and progressive KF in adults in our setup.

Key Words: Adults; Columbia classification; Focal segmental glomerulosclerosis; Histological variants; Kidney failure; Kidney failure with replacement therapy

©The Author(s) 2024. Published by Baishideng Publishing Group Inc. All rights reserved.

Core Tip: Focal segmental glomerulosclerosis (FSGS) is one of the most common glomerular diseases and a leading cause of kidney failure. FSGS is a heterogeneous disorder with many causes, varying pathogenesis and clinical courses. Columbia classification identified five histological variants of FSGS. The prognostic significance of these has remained controversial. Early studies found a good correlation of the variants with clinical presentation, treatment responses, and final outcomes. However, a more recent Japanese study found no prognostic value of the variants. The present study aimed to determine the clinical significance of these variants in a large sample of the Pakistani adult FSGS population.

Citation: Jafry NH, Manan S, Rashid R, Mubarak M. Clinicopathological features and medium-term outcomes of histologic variants of primary focal segmental glomerulosclerosis in adults: A retrospective study. *World J Nephrol* 2024; 13(1): 88028

URL: <https://www.wjgnet.com/2220-6124/full/v13/i1/88028.htm>

DOI: <https://dx.doi.org/10.5527/wjn.v13.i1.88028>

INTRODUCTION

Focal segmental glomerulosclerosis (FSGS) is a histological pattern of glomerular injury rather than a specific diagnosis. It can occur either in a primary or idiopathic form or may be associated with various systemic diseases, including autoimmune diseases, infections, drugs, and structural kidney diseases. The key pathological finding of FSGS on light microscopy is the obliteration involving a portion (segmental) of glomerular tufts of some (focal) but not all glomeruli by increased mesangial matrix. The underlying cause of idiopathic FSGS is uncertain although a putative circulating permeability factor is supposed to play a major role in its pathogenesis[1].

FSGS is one of the leading causes of glomerular diseases, particularly those presenting with nephrotic syndrome (NS), accounting for 20 to 40% of the pathological lesions in adult patients undergoing kidney biopsy for the evaluation of idiopathic NS[2-4]. It is also one of the most common glomerular diseases leading to kidney failure (KF) and KF requiring replacement therapy (KFRT)[2].

The classification of patients with FSGS is both challenging and controversial due to the wide variety of underlying etiologies, limited understanding of the pathogenesis, and the poor correlation between morphological lesions and response to treatment and clinical outcomes. The Columbia classification of FSGS provided a novel and pragmatic approach to classify FSGS based on histological features on kidney biopsy. This classification was supposed to help clinicians in the assessment of the prognosis of the disease and its response to therapy. The classification, first proposed by D'Agati *et al*[5], in 2004, envisioned five mutually exclusive histological variants of the disease, based entirely on light microscopic (LM) features[6]. These variants include collapsing (COL), not otherwise specified (NOS), tip (TIP), perihilar (PHI), and cellular (CEL) variants[2]. Since then, many studies of these variants have been conducted worldwide and have demonstrated a correlation of the variants with distinct clinical characteristics and prognostic and therapeutic outcomes[6-12]. The response rates are generally lowest for the COL variant, intermediate for the NOS variant, and highest for the TIP variant. On the other hand, the reported rates of KFRT are highest in the COL variant, intermediate in the NOS, and lowest in the TIP variant[13-16].

Other more recent studies have found no differences among these variants with respect to treatment responses and outcomes. A recent study from Japan by Kawaguchi *et al*[17] observed that FSGS variants alone have no significant impact on kidney outcomes after five years, while proteinuria remission was predictive of improved kidney prognosis irrespective of the variant. They suggested that specific strategies and interventions to achieve proteinuria remission for

each variant should be implemented for better kidney survival.

We previously reported that whatever the histological variant of FSGS, timely treatment with immunosuppressive drugs of all patients who fulfill the criteria is very important to achieve remission, either complete or partial remission (PR). Those patients who achieved remission did not progress to KF and did not require replacement therapy in the medium-term follow-up period[2,18].

As steroids and cyclosporine (CsA)-induced remission is associated with better long-term survival, it is important to study which type of FSGS variants are more likely to respond to steroids and CsA treatment and whether such treatment affects kidney survival[2,14,17].

This study aimed to determine the relative frequency, clinicopathologic presentations, and outcomes of histological variants of FSGS in our population.

MATERIALS AND METHODS

This retrospective observational study was conducted at the Department of Nephrology, Sindh Institute of Urology and Transplantation, Karachi, Pakistan. The study population comprised all adult patients (≥ 16 years) of either gender who were diagnosed with primary FSGS between January 1995 and December 2017. Secondary causes of FSGS were excluded. We did not analyze PHI and CEL variants in detail due to the small number of patients. Patients with missing information and irregular or erratic follow-up were also excluded.

All patients underwent ultrasound-guided percutaneous native kidney biopsies, which were processed and prepared according to standard guidelines, as described in detail in our previous studies[2,18,19]. The histological variants were diagnosed as per the criteria of the Columbia classification[5,6]. Briefly, the TIP variant was diagnosed when at least one segmental lesion involved the tip domain (outer 25% of the tuft next to the origin of the proximal tubule). It required the exclusion of COL and PHI variants. The tubular pole needs to be identified in the defining lesion. FSGS, NOS, was diagnosed when at least one glomerulus showed a segmental increase in the mesangial matrix obliterating the capillary lumina, with or without segmental glomerular capillary wall collapse but without overlying podocyte hyperplasia. It required the exclusion of PHI, CEL, TIP, and COL variants. The COL variant was diagnosed when at least one glomerulus showed segmental or global collapse and overlying podocyte hypertrophy and hyperplasia (Figure 1). The biopsies were reported by two experienced kidney pathologists and evaluated by LM, immunofluorescence, and electron microscopy. As the criteria for genetic testing in adult patients with FSGS are still unclear, genetic testing was not performed in this cohort of patients.

All patients with all variants of FSGS were treated in the same way. Briefly, unless contraindicated, all patients were treated with prednisolone, 1 mg/kg/d for the first six weeks followed by 0.75 mg/kg/d for the next six weeks. If no remission was achieved by the end of 12 wk, prednisolone was tapered over the next four weeks and stopped. If remission occurred at any time during treatment, the same dose of steroids was given for two more weeks before slow tapering. We did not employ different treatment protocols for different variants or different patients included in this study.

The steroid-resistant cases or those in which steroids were contraindicated were treated with CsA at a starting dose of 4 mg/kg/d. If a complete or PR occurred, CsA was continued for at least one year. If no response occurred by the end of two months, the use of CsA was discontinued.

Complete remission (CR) was defined as proteinuria ≤ 0.2 g/d or when the urine dipstick was negative for proteins with a stable serum creatinine concentration ($< 50\%$ increase from the baseline). PR was defined as proteinuria between 0.21-2.0 g/d with at least a 50% reduction in proteinuria from the baseline or albumin detected on dipstick (+1 to +4). Relapse was defined as proteinuria > 3 g/d after prior reduction of proteinuria to less than 2 g/d or albumin detected on dipstick (+1 to +3 or +4). Hypertension was diagnosed when patients were treated with antihypertensive drugs or with systolic blood pressure (BP) ≥ 140 mmHg or diastolic BP ≥ 90 mmHg. Elevated serum creatinine was defined as an increase of serum creatinine to > 1.4 mg/dL in male and > 1.2 mg/dL in female patients. KF was defined as a sustained increase of serum creatinine concentration $> 50\%$ from the baseline (at the time of kidney biopsy) or estimated glomerular filtration rate (eGFR) < 60 mL/min/1.73 m² according to the modification of diet in renal disease equation. KFRT was defined as the need for continuous dialysis or kidney transplantation.

Data were collected by reviewing the medical record files of all selected patients. Data were collected over the period from diagnosis of primary FSGS to the last follow-up. The demographic characteristics (age, gender), BP measurements, and laboratory investigations including proteinuria, serum albumin, serum creatinine, and urine dipstick results for albumin/red blood cells on initial and last visits were noted. Drug information and side effects of steroids were obtained. The outcome of all patients regarding sustained remission, or progression to KF/KFRT, and death was noted.

Data were entered and analyzed by SPSS software 22.0 (IBM Corp., Armonk, NY, United States). Descriptive statistics were used to summarize the continuous and categorical variables. Continuous variables such as age, BP, serum creatinine, serum albumin, 24-h proteinuria, and drug dosages, are presented as mean \pm SD and median \pm interquartile range (IQR), as appropriate. Categorical variables, *i.e.*, gender, sustained remission, KFRT, death, CR, PR, relapse, and hypertension are reported as frequencies and percentages. Mean differences for continuous variables and proportion differences for categorical variables between groups were compared using the student's *t*-test and Chi-square or Fisher's exact tests, as appropriate. Multivariate analysis for significant factors on univariate analysis was performed using the logistic regression method, while hazard ratios for risk factors of kidney outcome were calculated using the Cox regression model. Response rates, doubling of serum creatinine, and KFRT rates were compared between histological variants using ANOVA or Kruskal Wallis, and Chi-square tests as appropriate. A *P*-value ≤ 0.05 was considered statist-

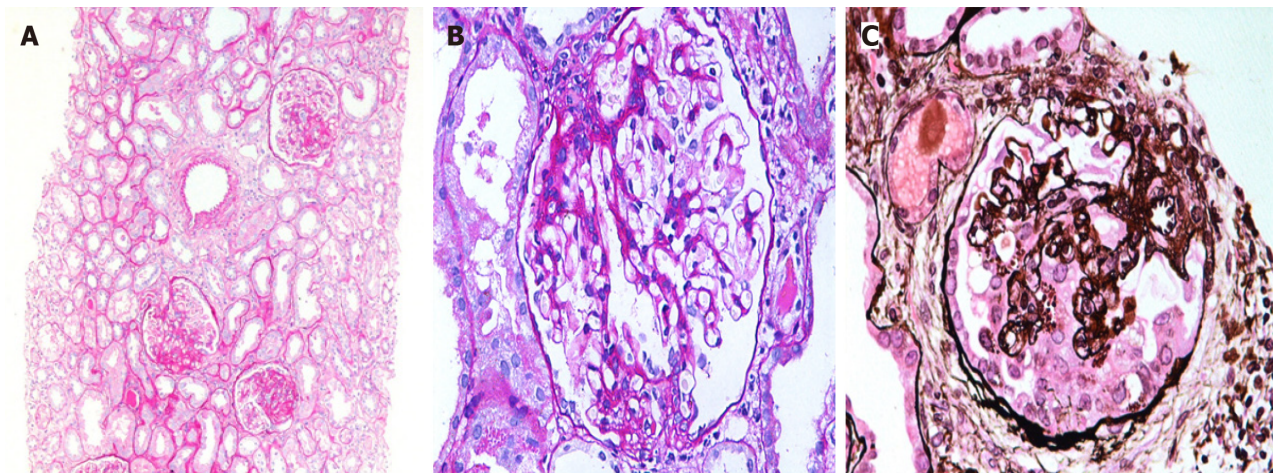


Figure 1 Histological features of common variants of focal segmental glomerulosclerosis. A: Medium-power view showing three glomeruli with segmental scars in indeterminate locations in a case of focal segmental glomerulosclerosis (FSGS), not otherwise specified. Mild patchy tubular atrophy and interstitial scarring is seen in the background [Periodic Acid-Schiff (PAS), × 200]; B: High-power view showing one glomerulus with segmental scar and adhesion formation involving the tip domain diagonally opposite the vascular pole, an example of the TIP variant (PAS, × 400); C: High-power view showing one glomerulus with segmental collapse of capillary tufts accompanied by podocyte hypertrophy and hyperplasia in a case of collapsing FSGS. Many proteinaceous droplets are also seen in the cytoplasm of podocytes (Jones's methenamine silver, × 400).

ically significant.

RESULTS

A total of 401 patients were diagnosed with primary FSGS during the study period. Among these, 352/401 (87.7%) had a designated Columbia histological variant. Among the latter, NOS was the commonest variant, found in 185 (53.9%) followed by the TIP variant in 100 (29.1%) patients. COL, CEL, and PHI variants were seen in 58 (16.9%), 6 (1.5%), and 3 (0.7%) patients, respectively. The three most common morphologic variants of FSGS (TIP, NOS, COL) comprised 343/352 (97.4%) and were included in the final analysis (Figure 2). The main demographic, clinical, and laboratory characteristics along with treatment information of these patients are shown in Table 1. There was no statistically significant difference in the mean ages of the patients with the three variants ($P = 0.7$). A statistically significant difference was observed in the diastolic BP where patients with the COL variant showed a higher value of 87.6 ± 14.1 mmHg compared to other histologic variants ($P = 0.04$). Initial proteinuria was nephrotic-range in all patients with a median (IQR) of 3774 (2216–5900) mg/24 h and there was no significant difference among the variants ($P = 0.418$). The initial eGFR was 83.9 (55.9 – 127.4) mL/min/ 1.73 m² in all patients with no significant difference among the three variants ($P = 0.463$). Elevated serum creatinine at presentation was found in 82 patients: Of these, 58 were males and 24 were females ($P = 0.25$). The mean follow-up duration in all patients was 36.5 ± 29.2 mo with no significant difference among the three variants ($P = 0.114$).

Of 343 patients, 302 (88%) received treatment with immunosuppressive agents combined with angiotensin-converting enzyme (ACE) inhibitors and angiotensin receptor blockers (ARBs) as shown in Figure 2. All patients received steroids as initial therapy as per our treatment protocol. The total duration of steroid therapy and total dose of steroids are shown in Table 1. The total dose of CsA is also shown in Table 1. The remaining patients were treated with ACE inhibitors or ARBs only due to reasons such as uncontrolled diabetes, intolerance of steroids, risk of osteoporosis, and other complications of immunosuppressive drugs.

Table 2 shows the details of pathological findings in the three morphological variants of FSGS. Upon review of the histopathological findings, a statistically significant difference was found among the three most common variants with respect to the number of glomeruli with global sclerosis ($P = 0.001$), number of glomeruli with segmental collapse ($P = 0.05$), and mild and moderate tubular atrophy ($P < 0.001$).

The treatment responses and clinical outcomes of all selected study subjects regarding complete and PR, or no remission, as well as progressive KF/KFRT, doubling of serum creatinine at last follow-up, final eGFR, relapse, and death of the patients are shown in Table 3. Of 302 patients, 84 (27.8%) achieved CR (Figure 2). Of these, 42 had the TIP variant, with a CR rate of 42/89 (47.2%), 40 had the NOS variant with a CR rate of 40/160 (25%), and 2 had the COL variant with a CR rate of 2/53 (3.7%) ($P < 0.001$). A total of 49/302 (16.2%) patients achieved PR. Among these, 11 had the TIP variant with a PR rate of 11/89 (12.4%), 27 had the NOS variant with a PR rate of 27/160 (16.8%), and 11 had the COL variant with a PR rate of 11/53 (20.8%) ($P = 0.005$). The highest percentage of no remission was found in patients with the COL variant at 40/53 (75.5%) followed by the NOS variant, 93/160 (58.1%) ($P < 0.001$) (Figure 3). The COL variant showed a marked decline in final eGFR and this was significant ($P < 0.001$) (Figure 4).

Among all treated patients, a doubling of serum creatinine was noted in 30 (9.9%) patients. Of these, 16/160 (10.0%) had the NOS variant, 1/89 (1.1%) had the TIP variant, and 13/53 (24.5%) had the COL variant ($P < 0.001$).

Table 1 Demographic, clinical, and laboratory parameters at the time of presentation of the three most common focal segmental glomerulosclerosis morphologic variants

Parameters	All patients (n = 343)	NOS (n = 185)	TIP (n = 100)	COL (n = 58)	P value
Age (yr), mean \pm SD	29.2 \pm 12.0	29.3 \pm 12.4	28.6 \pm 11.2	30.1 \pm 12.4	0.733
Male to female ratio	2.1:1	2.0:1	2.8:1	1.4:1	0.133
Systolic BP (mmHg), mean \pm SD	128.8 \pm 8.0	128.9 \pm 18.9	127.5 \pm 16.2	130.6 \pm 18.9	0.575
Diastolic BP (mmHg), mean \pm SD	84.6 \pm 12.6	84.8 \pm 12.0	82.5 \pm 12.5	87.6 \pm 14.1	0.045
Initial proteinuria (mg/24 h), IQR	3774 (2216-5900)	3602.5 (2125-5890)	4250 (2535-6530)	3900 (2218-4800)	0.418
Serum albumin (g/dL), mean \pm SD	2.0 \pm 1.2	2.0 \pm 1.3	2.2 \pm 0.9	2.1 \pm 1.1	0.863
Serum creatinine (mg/dL), mean \pm SD	1.1 \pm 0.9	1.1 \pm 0.7	1.1 \pm 1.1	1.2 \pm 0.7	0.830
Initial eGFR (mL/min/1.73 m ²), IQR	83.9 (55.9-127.4)	85.4 (57.8-128.3)	82.3 (59.5-125.9)	69.1 (41.9-127.3)	0.463
Elevated serum creatinine, n (%)	82 (23.9)	43 (52.4)	20 (24.4)	19 (23.2)	0.607
Males (> 1.4 mg/dL), n (%)	58 (70.7)	29 (67.4)	17 (85.0)	12 (63.2)	0.257
Females (> 1.2 mg/dL), n (%)	24 (29.3)	14 (32.6)	3 (15.0)	7 (36.8)	
Follow-up duration (month), mean \pm SD	36.5 \pm 29.2	39.1 \pm 31.5	31.4 \pm 24.9	37.0 \pm 27.7	0.114
Total steroid dose (mg), mean \pm SD	4282.1 \pm 1943.2	4266.9 \pm 1884.1	4366.7 \pm 2067.8	4186.1 \pm 1935.3	0.858
Duration of steroid treatment (wk), IQR	18 (14-23)	18 (14-23)	19 (15-23.5)	17 (13.5-20.5)	0.359
Total CsA dose (mg) mean \pm SD	27118.6 \pm 24904.8	30002.4 \pm 26537.5	25539.4 \pm 28050.2	21549.7 \pm 1657.7	0.447
Duration of CsA treatment (wk), IQR	16 (6.8-46.0)	16 (8.0-54.0)	17 (5.0-69.0)	12 (4.0-28.0)	0.727

NOS: Not otherwise specified; TIP: Tip; COL: Collapsing; CsA: Cyclosporine; BP: Blood pressure; eGFR: Estimated glomerular filtration rate; IQR: Interquartile range.

Table 2 Histopathology findings on kidney biopsies in the three most common focal segmental glomerulosclerosis morphologic variants

Histopathology	All patients (n = 343)	NOS (n = 185)	TIP (n = 100)	COL (n = 58)	P value
No. of glomeruli, mean \pm SD	17.3 \pm 8.4	16.4 \pm 8.7	18.1 \pm 6.8	17.3 \pm 9.7	0.370
No. of glomeruli with global sclerosis, mean \pm SD	2.6 \pm 2.2	2.1 \pm 2.3	2.0 \pm 0.8	3.6 \pm 2.5	0.001
No. of glomeruli with segmental collapse, mean \pm SD	2.9 \pm 2.7	2.4 \pm 1.8	2.6 \pm 2.8	13.7 \pm 3.3	0.050
Tubular atrophy, mild, n (%)	175/201 (87.1)	91 (52.0)	49 (28.0)	35 (20.0)	< 0.001
Tubular atrophy, moderate, n (%)	26/201 (12.9)	9 (36.6)	1 (3.8)	16 (61.5)	
Fibrointimal thickening of arteries, mild, n (%)	25/32 (98.1)	13 (52.0)	7 (28.0)	5 (20.0)	0.413
Fibrointimal thickening of arteries, moderate, n (%)	7/32 (21.9)	5 (27.8)	2 (22.2)	0 (0.0)	

NOS: Not otherwise specified; TIP: Tip; COL: Collapsing.

Regarding the development of KF, 23/53 (43.3%) patients with the COL variant developed progressive KF, 13/160 (18.1%) with the NOS variant, while none with the TIP variant progressed to KF during the mean follow-up period of 36.5 months (Figure 5). Similarly, 13/53 patients (24.5%) with the COL variant, 5/160 (3.1%) with the NOS variant, and none with the TIP variant required KFRT ($P < 0.001$). A total of four patients died, 1/53 (1.8%) with the COL variant, and 3/160 (1.8%) with the NOS variant with no significant difference among the three variants.

The final kidney and patient outcomes of the three common FSGS variants are shown in Figure 3. Patients with the TIP variant achieved remission more frequently (59.5%) than those with the NOS (41.8%), and COL (24.52%) variants ($P < 0.001$). The hazard ratio (HR) of complete response among patients with the COL variant was 0.163 (95%CI: 0.039-0.67) as compared to patients with the NOS variant. Patients with the TIP variant showed an HR of 2.50 (95%CI: 1.611-3.894) for CR compared to those with the NOS variant. Overall, progressive KF was observed more frequently in patients with the COL variant at 43.4% ($P < 0.001$). Among these, 24.53% of patients required kidney replacement therapy (KRT) ($P <$

Table 3 Responses to treatment and final clinical outcomes of focal segmental glomerulosclerosis morphologic variants in patients who received treatment, *n* (%)

Outcomes	All patients (<i>n</i> = 302)	NOS (<i>n</i> = 160)	TIP (<i>n</i> = 89)	COL (<i>n</i> = 53)	<i>P</i> value
Complete remission	84 (27.8)	40 (25.0)	42 (47.1)	2 (3.7)	< 0.001
Partial remission	49 (16.2)	27 (16.8)	11 (12.4)	11 (20.8)	0.005
No remission	169 (55.9)	93 (58.1)	36 (40.4)	40 (75.5)	< 0.001
Relapse	40 (13.2)	27 (16.9)	9 (10.1)	4 (7.5)	0.029
Final eGFR (mL/min/1.73 m ²), mean ± SD	92.1 ± 57.7	94.1 ± 59.1	110.3 ± 52.8	60.0 ± 46.9	< 0.001
Doubling of serum creatinine at last follow-up	30 (9.9)	16 (10.0)	1 (1.1)	13 (24.5)	< 0.001
Kidney failure	36 (11.19)	13 (8.1)	0 (0.0)	23 (43.3)	< 0.001
Hemodialysis	18 (5.9)	5 (3.1)	0 (0.0)	13 (24.5)	< 0.001
Expired	4 (1.3)	3 (1.8)	0 (0.0)	1 (1.8)	0.429

NOS: Not otherwise specified; TIP: Tip; COL: Collapsing; eGFR: Estimated glomerular filtration rate.

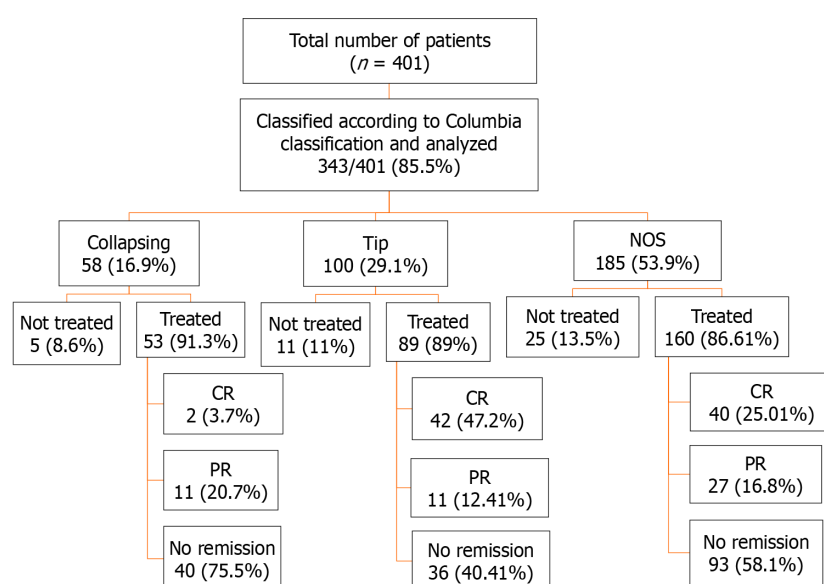


Figure 2 Flow diagram showing total number of patients and response rates of patients with the three histological variants of focal segmental glomerulosclerosis who were treated with steroids and cyclosporine. NOS: Not otherwise specified; CR: Complete remission; PR: Partial remission.

0.001). The HR of doubling of serum creatinine among patients with the COL variant was 14.577 (95%CI: 1.872-113.493) as compared to patients with the TIP variant.

DISCUSSION

This is the largest study with a cohort of 343 adult patients with biopsy-proven primary FSGS classified according to the Columbia classification from Pakistan. This study analyzed in detail the three most common morphological variants of primary FSGS, namely the NOS, TIP, and COL variants, as the numbers with the remaining two variants were very small. The vast majority of patients included in this study received treatment with steroids and CsA along with ACE inhibitors and ARBs. The total number of patients with the NOS variant was 185; of which, 160 (86.4%) patients received treatment. Likewise, there were 100 patients with the TIP variant; of which, 89 (89%) received treatment. There were 58 patients with the COL variant; of which, 53 (91.3%) patients received treatment. There is marked variation among these variants with regard to their frequency, clinical presentation, response to treatment, and prognosis in different regions of the world.

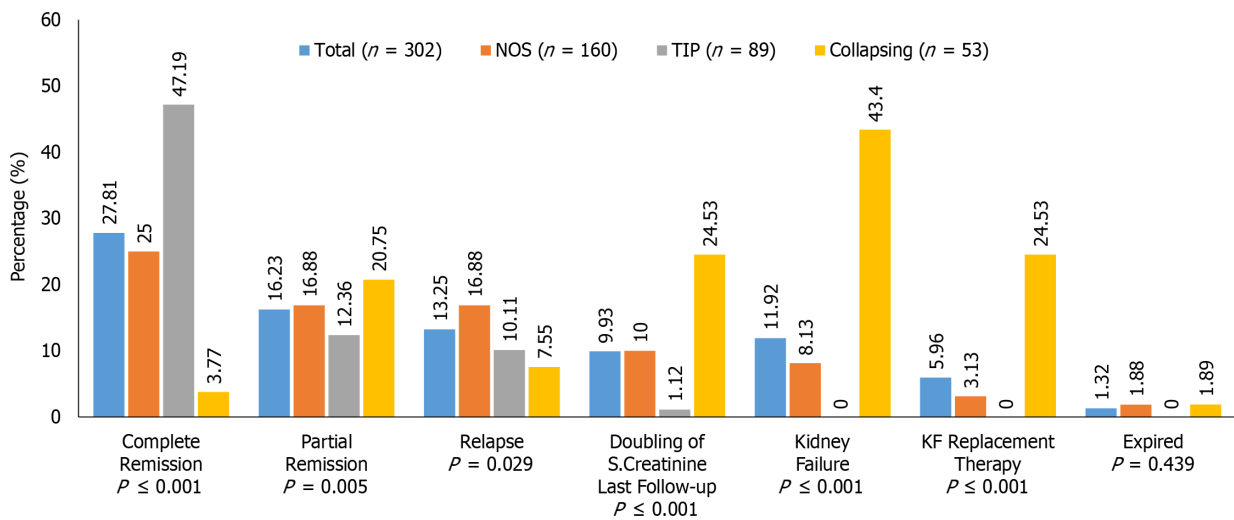


Figure 3 Treatment responses and final clinical outcomes of patients with the three most common histological variants of focal segmental glomerulosclerosis who were treated with immunosuppressive drugs ($n = 302$). NOS: Not otherwise specified; TIP: Tip; KF: Kidney failure.

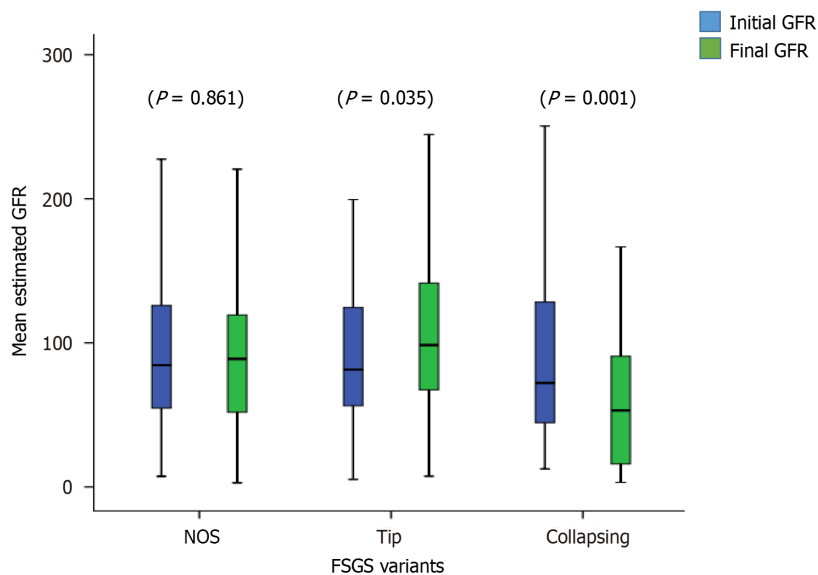


Figure 4 The means \pm SDs of the initial and final estimated glomerular filtration rates of patients with the three most common histological variants of focal segmental glomerulosclerosis. FSGS: Focal segmental glomerulosclerosis; GFR: Glomerular filtration rate; NOS: Not otherwise specified.

The exact reason for the paucity of PHI and CEL variants is not clear but there may be a misclassification of the CEL variant as the TIP variant, as CEL lesions can exist within the tuft at the tubular pole as the tip location and intracapillary expansive foam cells can be observed in both variants[7].

This is one of the largest cohorts of patients with COL FSGS in the Asian population as other studies from Asia did not include such a large number of patients with this lesion. A study on a Korean population of 111 patients with primary FSGS showed 63% NOS, 18% TIP, 15% PHI, 3% CEL, and only one patient with COL FSGS[8].

In general, there is a wide diversity in the prevalence of different variants of FSGS in different regions. In a Brazilian report, NOS was the most common variant, followed by COL[7]. It was also observed that there was a substantial overlap of criteria for the NOS and PHI variants as well as for COL and CEL variants. With regard to the CEL variant, it has been claimed that it is merely a form of the COL variant, and histologically it is very difficult to differentiate between these two variants. In fact, a common pathophysiological pathway affecting cell cycle regulatory proteins has been suggested for both variants[7]. A literature review also showed that the COL variant is less common in Whites than in the Black race as compared to NOS, TIP, PHI, and CEL variants, which are not common in the Black race[6,8,12].

A literature review of published studies on the Asian population showed that in China and India, TIP, NOS, and CEL variants are more frequent morphological variants with different treatment outcomes[16,20-22]. On the contrary, we found different results in our population. Shakeel *et al*[19] studied a large cohort of 184 patients from our center. They found that the COL variant was the second most common morphological variant of FSGS in our patients. These results

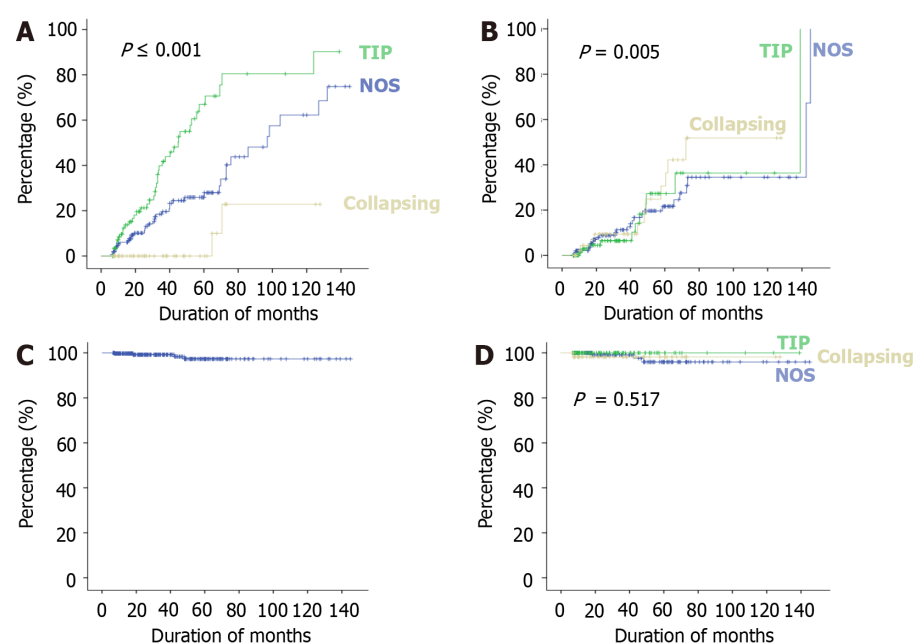


Figure 5 Treatment responses and cumulative kidney survival rates of patients with the three most common variants of focal segmental glomerulosclerosis. A: Kaplan-Meier plot of complete remission; B: Kaplan-Meier plot of partial remission; C: Kaplan-Meier plot of kidney survival (all patients); D: Kaplan-Meier plot of kidney survival by focal segmental glomerulosclerosis morphologic variants. NOS: Not otherwise specified; TIP: Tip.

were different to the Indian cohort as described previously. A more recent study from our institute evaluated the long-term outcome of adults with primary FSGS and showed a large number of patients with the COL variants after the NOS and TIP variants[2]. However, PHI and CEL variants were not common in our population as we did not find a significant number of patients with both these variants. A recent study from Japan showed a significant number of patients with PHI and CEL variants with no significant difference in outcome after treatment among different morphological variants of FSGS[17].

In our study, the majority of patients were young with an average age of 29 years, and all presented with nephrotic-range proteinuria. The mean follow-up period in our study was approximately 3 years. The primary outcome was the response to treatment and the secondary outcome was the composite outcome of the doubling of serum creatinine and KF with or without KRT. In addition, we also evaluated the improvement in eGFR as an additional outcome feature. We observed a good overall response to treatment in terms of achieving CR and PR. As in previous studies, this was highest in patients with the TIP variant, intermediate in those with the NOS variant, and lowest in patients with the COL variant. With the TIP variant, there was a > 34% increase in eGFR from the baseline after treatment which was a marked response to treatment, while in the NOS variant, eGFR increased to > 10% and in the COL variant, there was a decrease in eGFR of 13% at the end of the follow-up period. Only one patient with the TIP variant developed a doubling of serum creatinine and no patients required KFRT or expired in this group. There was a significant decline in eGFR in patients with the COL variant with only two patients out of 53 achieving CR, and 40 patients out of 53 achieved no remission. This poor outcome in those with the COL variant is similar to previous studies and observations worldwide[11,23-27]. This variant has always remained aggressive with a poor response to therapy[28-31]. However, a recent study from Japan showed almost similar outcomes for this variant compared to other variants except for the TIP variant. They proposed that this may be due to improved immunosuppressive treatment in recent years[17].

The poor outcomes in patients with the COL variant in our study are alarming in the sense that we have a large number of patients with the COL variant which ultimately can result in an increased burden of KFRT in patients in the much younger population. In those with the NOS variant, we observed that 93/160 (58.1%) patients achieved no remission. Overall, the TIP variant showed a very good response to therapy in terms of primary and secondary composite outcomes.

Two of our recent studies showed that almost half of the adults with primary FSGS achieved sustained remission with steroids and immunosuppressants, and consequently exhibited excellent short- to long-term kidney outcomes[2,18]. Almost the same results were obtained in this study in that of 302 patients, slightly less than half of the patients achieved sustained remission while 169 patients achieved no remission with a wide diversity of responses to treatment in different variants.

There are certain strengths as well as limitations in this study. The strengths include a large sample size, homogeneous race and uniform treatment protocol in all patients. Meticulous and accurate classification of morphological variants by experienced nephropathologists is also a strength of the study. We also noted some unique findings in this study. The prevalence of COL variant was quite high as compared with other regional studies. We also showed that morphological variants, if accurately classified, do have therapeutic and prognostic importance. The limitations include the single-center and retrospective nature of the study. The follow-up duration was not very long. Two variants were not analyzed due to very small numbers. Moreover, genetic testing was not performed in this cohort of patients, as currently the indications

for genetic testing in adult patients with FSGS are unclear. A pilot project to study the role of genetics in adult nephrotic patients at our center is in the pipeline and its results will be published in due course.

CONCLUSION

The histological variants of primary FSGS according to Columbia classification are associated with different clinicopathological presentations and are predictive of response to treatment and progressive KF. There was a large number of patients with the COL variant in this study, which is different to the rest of the Asian populations. These are results from a single center, and other studies are needed in order to compare the results and establish guidelines to effectively treat primary FSGS patients with different morphological variants.

ARTICLE HIGHLIGHTS

Research background

The classification of focal segmental glomerulosclerosis (FSGS) is controversial and challenging. There is still a lack of a unified and consensus-based approach to classify this disease, which will be both practical and clinically useful.

Research motivation

This study addressed the clinical utility of the morphological classification of FSGS in real-world scenarios. We aimed to investigate the therapeutic and prognostic significance of the morphological variants of FSGS in a large cohort of adult patients.

Research objectives

This study aimed to determine the relative prevalence, clinicopathologic presentations, and outcomes of the morphological variants of FSGS in a large cohort of adult patients at a single center in Pakistan.

Research methods

This retrospective study included all consecutive adults (≥ 16 years) with biopsy-proven primary FSGS from January 1995 to December 2017. Studied subjects were treated uniformly with steroids and cyclosporine. The response rates and kidney outcomes were compared between histological variants using appropriate statistical tests. Data were analyzed using SPSS version 22.0. A P -value ≤ 0.05 was considered statistically significant.

Research results

The not otherwise specified (NOS) variant was the most common, being found in 185 (53.9%) patients, followed by the TIP variant in 100 (29.1%) patients. Collapsing (COL), cellular, and perihilar variants were seen in 58 (16.9%), 6 (1.5%), and 3 (0.7%) patients, respectively. The response rates were highest in patients with the TIP variant and lowest in those with the COL variant. Kidney outcomes were best in patients with the TIP variant and worst in those with the COL variant. The NOS variant was intermediate.

Research conclusions

The morphological variants of FSGS are relevant and should be utilized to inform treatment and prognosis in individual patients. Combining these with other clinicopathological features to refine their predictive value needs to be investigated in future studies.

Research perspectives

A holistic approach to disease categorization needs to be developed, which is practical and clinical-friendly.

FOOTNOTES

Author contributions: Jafry NH and Mubarak M conceived and designed the study; Jafry NH, Manan S, Rashid R, and Mubarak M performed the research; all four participated in primary and final drafting; all read and approved the final manuscript; all four authors contributed significantly and equally to preparation of the manuscript.

Institutional review board statement: This study was reviewed and approved by the Ethics Committee of Sindh Institute of Urology and Transplantation.

Informed consent statement: All study participants or their legal guardian provided informed written consent regarding personal and medical data collection prior to study enrolment.

Conflict-of-interest statement: We have no financial relationships to disclose.

Data sharing statement: Data are available from the first author and can be shared on reasonable request.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <https://creativecommons.org/licenses/by-nc/4.0/>

Country/Territory of origin: Pakistan

ORCID number: Nazarul Hassan Jafrý 0000-0003-4800-9920; Shumaila Manan 0009-0005-1573-8459; Rahma Rashid 0000-0002-9332-2644; Muhammed Mubarak 0000-0001-6120-5884.

S-Editor: Qu XL

L-Editor: Webster JR

P-Editor: Zhao S

REFERENCES

- 1 Woo KT, Chan CM, Foo M, Lim C, Choo J, Chin YM, Teng EWL, Mok I, Kwek JL, Tan HZ, Loh AHL, Wong J, Kee T, Choong HL, Tan HK, Wong KS, Tan PH, Tan CS. Impact of focal segmental glomerulosclerosis over the past decade. *Clin Nephrol* 2023; **99**: 128-140 [PMID: 36633377 DOI: 10.5414/CN111022]
- 2 Jafrý N, Mubarak M, Rauf A, Rasheed F, Ahmed E. Clinical Course and Long-term Outcome of Adults with Primary Focal Segmental Glomerulosclerosis: A Retrospective Cohort Study. *Iran J Kidney Dis* 2022; **16**: 195-202 [PMID: 35714214]
- 3 Kazi JI, Mubarak M, Ahmed E, Akhter F, Naqvi SA, Rizvi SA. Spectrum of glomerulonephritides in adults with nephrotic syndrome in Pakistan. *Clin Exp Nephrol* 2009; **13**: 38-43 [PMID: 18685922 DOI: 10.1007/s10157-008-0075-0]
- 4 Nada R, Kharbanda JK, Bhatti A, Minz RW, Sakhuja V, Joshi K. Primary focal segmental glomerulosclerosis in adults: is the Indian cohort different? *Nephrol Dial Transplant* 2009; **24**: 3701-3707 [PMID: 19589787 DOI: 10.1093/ndt/gfp328]
- 5 D'Agati VD, Fogo AB, Bruijn JA, Jennette JC. Pathologic classification of focal segmental glomerulosclerosis: a working proposal. *Am J Kidney Dis* 2004; **43**: 368-382 [PMID: 14750104 DOI: 10.1053/j.ajkd.2003.10.024]
- 6 D'Agati V. Pathologic classification of focal segmental glomerulosclerosis. *Semin Nephrol* 2003; **23**: 117-134 [PMID: 12704572 DOI: 10.1053/snep.2003.50012]
- 7 Testagrossa LA, Malheiros DMAC. Study of the morphologic variants of focal segmental glomerulosclerosis: a Brazilian report. *J Bras Patol Med Lab* 2012; **48**: 211-215 [DOI: 10.1590/S1676-24442012000300009]
- 8 Kwon YE, Han SH, Kie JH, An SY, Kim YL, Park KS, Nam KH, Leem AY, Oh HJ, Park JT, Chang TI, Kang EW, Kang SW, Choi KH, Lim BJ, Jeong HJ, Yoo TH. Clinical features and outcomes of focal segmental glomerulosclerosis pathologic variants in Korean adult patients. *BMC Nephrol* 2014; **15**: 52 [PMID: 24666814 DOI: 10.1186/1471-2369-15-52]
- 9 Stokes MB, D'Agati VD. Morphologic variants of focal segmental glomerulosclerosis and their significance. *Adv Chronic Kidney Dis* 2014; **21**: 400-407 [PMID: 25168828 DOI: 10.1053/j.ackd.2014.02.010]
- 10 D'Agati VD, Alster JM, Jennette JC, Thomas DB, Pullman J, Savino DA, Cohen AH, Gipson DS, Gassman JJ, Radeva MK, Moxey-Mims MM, Friedman AL, Kaskel FJ, Trachtman H, Alpers CE, Fogo AB, Greene TH, Nast CC. Association of histologic variants in FSGS clinical trial with presenting features and outcomes. *Clin J Am Soc Nephrol* 2013; **8**: 399-406 [PMID: 23220425 DOI: 10.2215/CJN.06100612]
- 11 Deegens JK, Steenberg EJ, Borm GF, Wetzels JF. Pathological variants of focal segmental glomerulosclerosis in an adult Dutch population-epidemiology and outcome. *Nephrol Dial Transplant* 2008; **23**: 186-192 [PMID: 17704112 DOI: 10.1093/ndt/gfm523]
- 12 Chun MJ, Korbet SM, Schwartz MM, Lewis EJ. Focal segmental glomerulosclerosis in nephrotic adults: presentation, prognosis, and response to therapy of the histologic variants. *J Am Soc Nephrol* 2004; **15**: 2169-2177 [PMID: 15284302 DOI: 10.1097/01.ASN.0000135051.62500.97]
- 13 Arias LF, Jiménez CA, Arroyave MJ. Histologic variants of primary focal segmental glomerulosclerosis: presentation and outcome. *J Bras Nefrol* 2013; **35**: 112-119 [PMID: 23812568 DOI: 10.5935/0101-2800.20130019]
- 14 Korbet SM. Treatment of primary FSGS in adults. *J Am Soc Nephrol* 2012; **23**: 1769-1776 [PMID: 22997260 DOI: 10.1681/ASN.2012040389]
- 15 Thomas DB, Franceschini N, Hogan SL, Ten Holder S, Jennette CE, Falk RJ, Jennette JC. Clinical and pathologic characteristics of focal segmental glomerulosclerosis pathologic variants. *Kidney Int* 2006; **69**: 920-926 [PMID: 16518352 DOI: 10.1038/sj.ki.5000160]
- 16 Tang X, Xu F, Chen DM, Zeng CH, Liu ZH. The clinical course and long-term outcome of primary focal segmental glomerulosclerosis in Chinese adults. *Clin Nephrol* 2013; **80**: 130-139 [PMID: 23380386 DOI: 10.5414/CN107607]
- 17 Kawaguchi T, Imasawa T, Kadamura M, Kitamura H, Maruyama S, Ozeki T, Katafuchi R, Oka K, Isaka Y, Yokoyama H, Sugiyama H, Sato H. Focal segmental glomerulosclerosis histologic variants and renal outcomes based on nephrotic syndrome, immunosuppression and proteinuria remission. *Nephrol Dial Transplant* 2022; **37**: 1679-1690 [PMID: 34499164 DOI: 10.1093/ndt/gfab267]
- 18 Jafrý N, Ahmed E, Mubarak M, Kazi J, Akhter F. Raised serum creatinine at presentation does not adversely affect steroid response in primary focal segmental glomerulosclerosis in adults. *Nephrol Dial Transplant* 2012; **27**: 1101-1106 [PMID: 21804085 DOI: 10.1093/ndt/gfr430]
- 19 Shakeel S, Mubarak M, I Kazi J, Jafrý N, Ahmed E. Frequency and clinicopathological characteristics of variants of primary focal segmental glomerulosclerosis in adults presenting with nephrotic syndrome. *J Nephropathol* 2013; **2**: 28-35 [PMID: 24475423 DOI: 10.5812/nephropathol.8959]
- 20 Das P, Sharma A, Gupta R, Agarwal SK, Bagga A, Dinda AK. Histomorphological classification of focal segmental glomerulosclerosis: a critical evaluation of clinical, histologic and morphometric features. *Saudi J Kidney Dis Transpl* 2012; **23**: 1008-1016 [PMID: 22982914 DOI: 10.4103/1319-2442.100883]
- 21 Nuguri S, Swain M, Padua M, Gowrishankar S. A Study of Focal and Segmental Glomerulosclerosis according to the Columbia Classification

- and Its Correlation with the Clinical Outcome. *J Lab Physicians* 2023; **15**: 431-436 [PMID: 37564237 DOI: 10.1055/s-0043-1761930]
- 22 **Swarnalatha G**, Ram R, Ismal KM, Vali S, Sahay M, Dakshinamurthy KV. Focal and segmental glomerulosclerosis: does prognosis vary with the variants? *Saudi J Kidney Dis Transpl* 2015; **26**: 173-181 [PMID: 25579744 DOI: 10.4103/1319-2442.148772]
 - 23 **Schwartz MM**, Korbet SM, Rydell J, Borok R, Genchi R. Primary focal segmental glomerular sclerosis in adults: prognostic value of histologic variants. *Am J Kidney Dis* 1995; **25**: 845-852 [PMID: 7771480 DOI: 10.1016/0272-6386(95)90566-9]
 - 24 **Carney EF**. Glomerular disease: association of FSGS histologic variants with patient outcomes. *Nat Rev Nephrol* 2013; **9**: 65 [PMID: 23296292 DOI: 10.1038/nmeph.2012.272]
 - 25 **Choi MJ**. Histologic classification of FSGS: does form delineate function? *Clin J Am Soc Nephrol* 2013; **8**: 344-346 [PMID: 23430205 DOI: 10.2215/CJN.00660113]
 - 26 **Tsushima A**, Matsukuma Y, Ueki K, Tanaka S, Masutani K, Nakagawa K, Mitsui K, Uesugi N, Katafuchi R, Tsuruya K, Nakano T, Kitazono T. Utility of Columbia classification in focal segmental glomerulosclerosis: renal prognosis and treatment response among the pathological variants. *Nephrol Dial Transplant* 2020; **35**: 1219-1227 [PMID: 30649467 DOI: 10.1093/ndt/gfy374]
 - 27 **Trivedi M**, Pasari A, Chowdhury AR, Abraham-Kurien A, Pandey R. The Spectrum of Focal Segmental Glomerulosclerosis from Eastern India: Is It Different? *Indian J Nephrol* 2018; **28**: 215-219 [PMID: 29962672 DOI: 10.4103/ijn.IJN_115_17]
 - 28 **Han MH**, Kim YJ. Practical Application of Columbia Classification for Focal Segmental Glomerulosclerosis. *Biomed Res Int* 2016; **2016**: 9375753 [PMID: 27247945 DOI: 10.1155/2016/9375753]
 - 29 **Ozeki T**, Nagata M, Katsuno T, Inagaki K, Goto K, Kato S, Yasuda Y, Tsuboi N, Maruyama S. Nephrotic syndrome with focal segmental glomerular lesions unclassified by Columbia classification; Pathology and clinical implication. *PLoS One* 2021; **16**: e0244677 [PMID: 33400710 DOI: 10.1371/journal.pone.0244677]
 - 30 **Meehan SM**, Chang A, Gibson IW, Kim L, Kambham N, Laszik Z. A study of interobserver reproducibility of morphologic lesions of focal segmental glomerulosclerosis. *Virchows Arch* 2013; **462**: 229-237 [PMID: 23262784 DOI: 10.1007/s00428-012-1355-3]
 - 31 **Vlasić-Matas J**, Glavina Durdov M, Capkun V, Galesić K. Prognostic value of clinical, laboratory, and morphological factors in patients with primary focal segmental glomerulosclerosis - distribution of pathological variants in the Croatian population. *Med Sci Monit* 2009; **15**: PH121-PH128 [PMID: 19789521]



Published by **Baishideng Publishing Group Inc**
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

Telephone: +1-925-3991568

E-mail: office@baishideng.com

Help Desk: <https://www.f6publishing.com/helpdesk>

<https://www.wjgnet.com>

