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An approximation to the prevalence of focal segmental glomerulosclerosis: A systematic review of world literature over the past 32 years.

Short title: Focal Segmental Glomerulosclerosis in the world.

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Abstract

Background: Focal segmental glomerulosclerosis (FSGS) is a histopathological lesion characterized by scarring in specific sections of some glomeruli, accompanied by podocyte injury. Worldwide, the prevalence of FSGS and its temporal trends have not been sufficiently studied. However, some reports suggest an increase in the frequency of FSGS in recent decades. Understanding the epidemiology of FSGS is crucial for clinicians to improve diagnosis and treatment. **Objective:** This study critically evaluates global prevalence trends of FSGS over the past 32 years (1992-2024), highlighting variations between countries through a systematic review. Methods: A systematic search of Medline, Embase and ScienceDirect was conducted to identify relevant studies. The reliability of prevalence data was assessed by critical appraisal of selected publications. Results: The prevalence of FSGS varies significantly between regions. East Asian countries have a relatively low prevalence, with a mean around 7%. In contrast, countries in South Asia, the Middle East and the Americas have a higher prevalence of around 18%. European countries show an intermediate prevalence of about 11%. African countries do not show a clear pattern, with high and low prevalence rates in different countries. Conclusions: The prevalence of FSGS differs by geographic region and ethnicity. While South Asian countries have maintained a consistently low prevalence, other regions have experienced an increase in FSGS cases over time. This study improves the understanding of global patterns of FSGS, providing valuable epidemiological insights for clinicians and researchers.

Keywords: Focal segmental glomerulosclerosis, glomerulonephritis, kidney biopsy, prevalence.

1. Introduction

Focal segmental glomerulosclerosis (FSGS) is a histopathological lesion characterized by scarring (sclerosis) affecting specific sections (segmental) of some glomeruli (focal), accompanied by podocyte injury ¹. Although previously considered a single disease, FSGS is now recognized as a heterogeneous condition with diverse etiologies, clinical presentations and responses to treatment. The defining characteristic of FSGS is podocyte injury and loss, which can occur as a primary disorder or as a secondary response to various glomerular stressors. As the disease progresses, localized segmental sclerosis can spread to involve entire glomeruli, eventually culminating in complete glomerular dysfunction ¹.

Since the 1970s, FSGS has attracted increasing attention due to its role as a leading cause of end-stage renal disease (ESRD) worldwide ².

The morphological classification of FSGS, based on the Columbia system ³, classified the disease into five variants: collapsing, tip, cellular, perihilar, and not otherwise specified. However, relying solely on this framework to make therapeutic decisions proved to be insufficient, as it did not take into account key genetic factors or syndromic presentations, often leading to ineffective treatment. In response, a more comprehensive approach has emerged that considers both clinical and pathological features, redefining FSGS into primary, secondary, genetic and indeterminate forms ⁴. Primary FSGS is thought to be caused by an unknown circulating factor, in the absence of an identifiable underlying cause. In recent years, several studies have identified anti-nephrin antibodies in a substantial proportion of patients with primary FSGS, suggesting their potential role in the disease ⁵. These antibodies have been proposed as candidate permeability factors; however, their direct involvement in the pathophysiology of FSGS has not yet been conclusively demonstrated. Other proposed candidates include soluble urokinase plasminogen activator surface receptor (suPAR), apolipoprotein A1b (APOA1b), cardiotrophin-like cytokine factor 1 (CLCF1), and anti-CD40 antibodies ⁶. In contrast, secondary FSGS develops as a consequence of systemic conditions or external influences, such as viral infections, exposure to toxins or drugs, and glomerular hyperfiltration due to factors such as obesity, congenital renal anomalies, solitary kidney or reflux nephropathy ⁴. Genetic FSGS, on the other hand, is the result of mutations in genes essential for podocyte function and structural integrity, leading to progressive renal damage 7.

Although it is recognized that the overall incidence of FSGS has increased ⁸, its prevalence varies significantly depending on factors such as geographic region, study period, race, age, and sex distribution. The objective of this review is to critically evaluate prevalence studies published in the last 32 years, providing valuable information on trends, regional differences, and changes in disease rates. By conducting a systematic analysis of the global prevalence and incidence of FSGS, we aim to improve understanding of its epidemiological patterns and suggest future research and health care strategies.

2. Methods

The search strategy followed the preferred reporting items for systematic reviews and metaanalysis (PRISMA) guidelines. A comprehensive search of different information sources was carried out in Medline, Embase, and ScienceDirect databases (1992-2024) was performed using the terms "glomerulonephritis", "nephropathy", "kidney disease", "prevalence", "incidence", "epidemiology", "focal segmental glomerulosclerosis", and/or "focal segmental glomerulosclerosis epidemiology", The eligibility criteria for the included studies required them to present original research, were published in English or Spanish, and contained relevant FSGS epidemiological information about trends, epidemiology or incidence of FSGS. Each record was independently reviewed by two researchers based on its title and abstract. In cases of disagreement regarding a study's inclusion, a third researcher evaluated the record and made the final decision. All studies were considered and reviewed from 10 January 2023 to 30 December 2024, regardless of their focus or age group, as long as they contained information according to the inclusion criteria. Exclusion criteria included duplicate publications, insufficient information on incidence, unavailability of full text or incomplete data, and reporting bias. Review articles, transplant registries, conference abstracts, unpublished manuscripts, small sample size (less than 100 patients) in countries with multiple reports and studies on recurrent FSGS were also excluded. Only descriptive statistics were used from the original article. Alternative meta-analyses or advanced statistical models were not performed due to several limitations inherent to the study. These include the use of registry data that may not accurately reflect the general population in some countries, discrepancies in the time periods covered by the various registries, and significant heterogeneity in renal biopsy practices across countries. This variability includes differences

in clinical indications for biopsy, procedural strategies, and diagnostic criteria, all of which limit the comparability of the data. We evaluated the language of publication biases, but only 8 articles were discarded as being written in Hebrew, Italian, French, Croatian, German, Chinese, Hungarian and Romanian. Titles and abstracts were reviewed by the authors, and selected studies were evaluated in more detail. We considered frequency and prevalence data reported by the authors, incorporating information from national biopsy registries, case reports, and case series. This work was not registered in PROSPERO because the project had already commenced before registration was considered, and it was later decided to proceed without registration given the stage of progress at that time.

3. Results

The initial literature search identified 1,047 articles, which were systematically selected according to predefined inclusion and exclusion criteria. After this screening process, only 130 original studies met the analysis criteria. A detailed breakdown of the screening process, including the number of articles included and excluded at each stage, is presented in Figure 1. Across all continents, Asia was the continent with the highest number of reports, 67 in total. Among them, China contributed the largest number of studies with 16, followed by India with 10, Japan with 7 and Pakistan with 4. South Korea and Iran provided 4 studies, while Saudi Arabia, Turkey and Nepal contributed 3 each. Lebanon and Jordan submitted 2 reports, while the United Arab Emirates, Taiwan, Thailand, Oman, Iraq, Bangladesh and Sri Lanka, Kuwait and Singapore contributed 1 study each. Europe provided 33 reports, with Italy leading with 5, followed by Romania with 3, Poland, Spain, the Czech Republic, Denmark, Serbia, Belgium, Germany and Croatia with 2 each. Sweden, France, Cyprus, Lithuania, Estonia, Portugal, Finland, Norway and United Kingdom received 1 report each. The Americas contributed 22 reports, with Brazil leading the way with six studies, followed by the United States and Colombia with five each, Mexico with 2 and Chile, Uruguay, Peru and Canada with 1 each. Africa presented 9 reports, 4 from Nigeria and 1 each from Egypt, South Africa, Morocco, Chad and Sudan. Oceania was the continent with the fewest reports, with only 3, 2 from Australia and 1 from New Caledonia. **Table 1** provides an overview of the main characteristics of the studies included in this analysis, and Supplementary Table 1 contains the complete list of studies.

3.1. Asia

In Asia, the prevalence of FSGS varies significantly from region to region. In general, South Asian countries, such as India, Pakistan and Sri Lanka, together with Middle Eastern countries, such as Iran, Jordan, Saudi Arabia, Iraq, Lebanon, United Arab Emirates and Oman, have a high prevalence of FSGS, with a mean of about 19%, although some countries, such as Bangladesh and Nepal, show more moderate levels. In contrast, East Asian countries, including China, Korea, Japan and Thailand, tend to have a lower prevalence, with an average of approximately 6.5%, with some exceptions, such as Taiwan, where the prevalence is slightly higher.

3.1.1. China

The 20 reports from China span a study period from 1979 to 2020, encompassing data from several medical institutions, with some temporal overlap between studies. Together, these studies analyzed a total of 181,384 samples, with individual sample sizes ranging from 162 to 62,569. The reported prevalence of FSGS remained relatively low, with a mean of 7.01%, and values ranging from 2.45% to 21.6%. Despite this variability, prevalence rates were generally consistent across studies. Based on five studies conducted in the adult population, the prevalence of FSGS has shown slight variations over time. From 1993 to 1997, it was estimated at approximately 3.48%, increasing to 5.47% from 1998 to 2002. From 2003 to 2019, the prevalence remained relatively stable at around 4.53%. Distinguishing differences in prevalence between children and adults is challenging, as many studies analyze both groups together. However, based on data from three studies, the estimated prevalence of FSGS in children (≤14 years old) is approximately 4.67%. In contrast, adults (>14 years old), for whom more reports are available, show a slightly higher mean prevalence of approximately 7.06%.

3.1.2. India

Studies conducted in India covered the period from 1990 to 2020, with some temporal overlap between studies, and reported a higher prevalence of FSGS compared to those in China. Across all studies, the overall mean prevalence was 18.26% among the 8,325 biopsies analyzed, with values ranging from 8.02% to 22.58%. The number of samples analyzed per study ranged from 65 to 3,257. When looking at specific age groups, six studies focusing on

the pediatric population (≤14 years old) estimated a prevalence of approximately 14.53%, while eight studies in adults reported a slightly higher prevalence of around 17.05%.

3.1.3. Japan

The studies conducted in Japan covered the period from 2007 to 2021, with some temporal overlap and the inclusion of data from multiple medical institutions, including some large-scale studies. According to these studies, the mean incidence of FSGS was 6.36%, based on an analysis of 49,171 biopsies. The number of samples analyzed per study ranged from 438 to 32,254, with a median of 2,802. The overall variation in prevalence was small, ranging from 3.5% to 10.85%. Only one study focused on the pediatric population, reporting a prevalence of 3.7% during the period 2007 to 2017. In contrast, the adult population was analyzed in seven studies, with a mean prevalence of 6.24%.

3.1.4. Pakistan

The six studies conducted in Pakistan analyzed a total of 1,375 kidney biopsies over a period from 1997 to 2021. Most reports focus on the pediatric population (\leq 14 years of age) and reported a high prevalence of FSGS, with a mean of 25.54% and a range between 12% and 40.4%. Only one study examined the adult population and reported an even higher prevalence of 30.86%.

3.1.5. South Korea

The South Korean population studies spanned from 1973 to 2013 and analyzed a total of 6,657 biopsy samples. Like China and Japan, the prevalence of FSGS among the Korean population is generally low, at 6.21%. However, this rate varies by age group. In pediatric patients (≤14 years of age), one study reports a prevalence of 4%. Among adults under 65 years of age, two studies estimate it at 5.15%, while among those over 65, the prevalence increases to 12.2%.

3.1.6. Iran

Similar to India and Pakistan, the prevalence of FSGS in adult Iranian population is relatively high, averaging 21.63% based on three studies. These studies, conducted between 2006 and 2018, analyzed a total of 4,803 biopsy samples.

3.1.7. Saudi Arabia

In Saudi Arabia, three studies conducted between 1989 and 2020 reported an average FSGS prevalence of 19.3% based on an analysis of 940 biopsy samples.

3.1.8. Turkey

In Turkey, the mean prevalence of FSGS was approximately 15.22%, based on an analysis of 8,381 biopsies in three studies. Two of these studies focused on the pediatric population and reported a mean incidence of 11.88%, while the third study, conducted in adults, found a prevalence of approximately 21.9%.

3.1.9. Jordan

In Jordan, the mean prevalence of FSGS is estimated at 18.44% according to three studies. However, the total sample size was relatively small, with only 260 biopsies analyzed between 2006 and 2020. Two studies reported a prevalence of 22% in the pediatric population, while one study in adults found a mean prevalence of 11.32%.

3.1.10. Nepal

In Nepal, the prevalence was moderate, estimated at approximately 14.85% based on three studies conducted between 2001 and 2023. However, this finding is based on a limited sample size, with only 549 biopsies analyzed.

3.2. Europe

Overall, the prevalence of FSGS in Europe appears to be relatively constant in all countries for which data were available (Italy, Romania, Poland, Spain, Sweden, Czech Republic, Denmark, Serbia, Belgium, France, Germany, Cyprus, Lithuania, Estonia, Croatia and the United Kingdom). In all these regions, prevalence remains moderate, with an average of around 10%.

3.2.1. Italy

The mean prevalence of FSGS in Italy was about 13.57%, based on 38,228 biopsies analyzed in seven studies conducted between 1970 and 2010, involving multiple medical institutions with some overlap between studies. Based on two studies in pediatric patients, the prevalence in this population was about 10.05%, while based on five studies in adult patients, it was slightly higher at 14.08%. Over time, the prevalence has gradually increased, as indicated by two studies: from 5.2% between 1907 and 1974, to 6.3% between 1975 and 1979, to 6.7% between 1980 and 1984, to 9% between 1985 and 1989, and to 11.2% between 1990 and 1994.

3.2.2. Romania

In Romania, the mean prevalence of FSGS was approximately 9.3%, according to four reports covering the period from 1995 to 2023, in which a total of 2,127 biopsies were analyzed. Incidence rates varied among the studies, with one reporting an incidence of 0.7 cases per million people per year and another reporting 10 cases per million people per year.

3.3. The Americas

Almost all countries studied in the Americas, including the United States, Brazil, Colombia, Mexico, Uruguay, Peru and Canada, report a high prevalence of FSGS, with some exceptions, such as Chile. Notably, Brazil is the country with the highest prevalence in the region.

3.3.1. Brazil

In Brazil, the prevalence of FSGS was remarkably high, ranking as the second highest among all the countries studied. However, Brazil had a much larger sample size than Nigeria, which reported the highest prevalence. A prevalence of approximately 33.13% was found in 15,666 biopsies analyzed in six studies conducted between 1992 and 2018.

3.3.2. United States

In the United States, the prevalence of FSGS was relatively high, with a mean of 20.13% according to six studies analyzing 5,173 biopsies collected between 1974 and 2020, with some overlap between studies. The incidence was recorded at approximately 1.6 cases per 100,000 people per year according to two studies, with an increase observed when comparing the periods 1994 to 2003 and 2004 to 2013.

3.3.3. Colombia

In Colombia, the estimated prevalence of FSGS was high, around 20.39%, based on 12,354 biopsies analyzed in four studies conducted between 1998 and 2017. Among the pediatric population (\leq 15 years old), the prevalence was estimated at 21.9%, while in the adult population it was higher, at 25.17%, based on three studies each.

3.4. Africa

In African countries, prevalence varies from moderate, around 11% in South Africa, Morocco, Chad and Sudan, to high, above 20% in Nigeria and Egypt. However, these estimates are based on a limited number of samples tested.

3.4.1. Nigeria

Nigeria had the highest prevalence of all the countries studied, with a mean of 33.84%. However, this finding is based on a relatively small sample, with only 360 biopsies analyzed in five studies conducted between 1997 and 2022. Notably, most of these studies focused on children, and only one included both children and adults.

3.5. Oceania

In Oceania, the prevalence of FSGS is relatively high, averaging approximately 18.53%. This estimate is based on an analysis of 4,531 samples collected in Australia and New Caledonia between 1982 and 2011.

4. Discussion

The prevalence of FSGS varies significantly from region to region (**Figure 2**). In general, in Asia, South Asian and Middle Eastern countries have a high prevalence, with an average of about 19%, while East Asian countries have lower rates, with an average of 6.5%. In Europe, the prevalence of FSGS remains relatively uniform across countries, with an average of 10%. In the Americas, most countries have a high prevalence. Africa shows a spectrum of prevalence, ranging from moderate to high rates exceeding 20%, but data are limited. A systematic review and meta-analysis in African populations reported a pooled FSGS prevalence of 26.1%. Prevalence rates were similarly high across sub-Saharan regions [West (34.9%), East (33.5%), and South (34.8%)], while notably lower in North Africa (19.8%) ⁹. Oceania has a relatively high average prevalence of 18.5%.

It is important to note that data from many countries are based on limited samples, not numerically representative of the population, which means that actual prevalence rates of FSGS may differ from those reported. In addition, the reported incidence may vary according to the type of healthcare facility, such as primary, secondary or tertiary level hospitals, which may directly influence the observed prevalence of primary and secondary glomerulopathies. Despite these limitations, this study provides valuable information on an estimate of the overall prevalence of FSGS in different countries and, in some cases, highlights trends and changes over time (**Figure 3**).

The prevalence of FSGS is influenced by a complex interaction of genetic, dietary, environmental and health factors. An example of this is that in African countries with a high prevalence of FSGS, such as Nigeria and Egypt, a high presence of high-risk variants of the APOL1 gene has been identified ¹⁰. In contrast, populations in East Asian countries such as

China, Japan and Korea have a lower prevalence of these high-risk APOL1 variants, which may partly explain the lower incidence of FSGS in these regions ¹¹. Dietary patterns also play a critical role. In Western countries, such as the United States and Brazil, diets high in sodium and protein are common. These dietary factors can exacerbate hypertension and hyperfiltration injury, contributing to the higher rates of FSGS observed in these populations ¹². In contrast, the diets of East Asian countries, rich in omega-3 fatty acids due to high fish consumption, have anti-inflammatory properties that may provide protective effects against glomerular damage ¹³. Similarly, the Mediterranean diet, predominant in Southern Europe, is associated with lower rates of cardiovascular and metabolic diseases, key risk factors for kidney disease 14. Additionally, rapid urbanization in countries like India and Brazil has led to lifestyle changes, including increased sedentary behaviors and dietary shifts, which can amplify metabolic risk factors for kidney disease. Notably, countries like Japan and Korea have maintained relatively low obesity rates, which could also contribute to their reduced prevalence of FSGS and other kidney-related conditions ¹⁵. It is important to note that disparities in access to genetic testing across countries may also contribute to the higher prevalence of FSGS observed in biopsy samples from lower-resource settings. In wealthier countries, genetic forms of glomerular disease (e.g., APOL1-associated nephropathy) are more often diagnosed through genetic testing, potentially reducing the need for renal biopsy in these cases. As a result, such cases may be underrepresented in biopsy registries from highresource countries and overrepresented in those from regions with limited access to genetic diagnostics.

Historical reports highlight the global variation in FSGS incidence over time. In 2011, McGrogan et al. documented an incidence of 0.2–1.1 cases per 100,000 person-years ¹⁶, while Jegatheesan et al. in 2016 reported an incidence of 1.02 cases per 100,000 person-years ¹⁷. In Germany, the incidence increased from 0.1 cases per 100,000 person-years (1990–1997) to 0.6 (2006–2013) ¹⁸. Similarly, Uruguay reported stable incidence rates across five time periods (1990–2014), ranging between 6.93 and 12.01 per million person-years ¹⁹. Italy also observed an increasing trend, with Schena et al. reporting an incidence of 2.3 per million population (pmp) in 1993 and FSGS accounting for 16.6% of nephrotic syndrome cases by 2004 ²⁰. Meanwhile, in India, prevalence has varied widely, ranging from 13.1% to 30.6% across different cohorts, reflecting diverse genetic and environmental influences.

Recent trends in FSGS prevalence and incidence reveal changing patterns over time. In North America, Molnár et al. (2023) observed a prevalence of 17.7%, noting a downward trend in the past three years ²¹. Conversely, Nakagawa et al. (2023) documented stabilization of FSGS incidence in Europe after a decline from 18.6 pmp in 2000 to 14.5 pmp in 2013 ²². In Latin America, countries like Colombia reported an increase in prevalence from 20.6% in 2013, while in Brazil, prevalence peaked at 55.4% between 2000–2005 before declining to 25.8% by 2018.

Age and gender significantly influence FSGS prevalence. Males are predominantly affected, with higher frequencies observed in pediatric populations, particularly among those with corticosteroid-resistant nephrotic syndrome. In contrast, geriatric populations exhibit much lower rates of this podocytopathy ⁶. A study in Uruguay spanning 25 years (1990–2014) showed FSGS was most prevalent among individuals aged 15–50 years, with declining prevalence in those over 65 years of age ¹⁹. Pediatric populations are especially affected, with prevalence rates ranging from 8.5–11.6% in Italy, 18.3–25.9% in Pakistan, and 28.7% in Colombia, while India reported 9.23%, with 36.2% of cases being steroid-resistant ^{23–27}. A systematic review on the epidemiology of childhood nephrotic syndrome in Africa reported a rise in the prevalence of FSGS from 14% in studies conducted before 1990 to 40% in those published after 1990. Regional differences were also observed, with the highest proportion of FSGS found in Central Africa (33%), followed by Southern (22.8%) and Northern Africa (21.7%). The lowest proportions were reported in Eastern (19.4%) and Western (19.1%) Africa ²⁸.

Despite the inherent limitations of data collection and representation, this analysis significantly improves our understanding of global patterns of FSGS. By offering detailed epidemiological insight, it provides clinicians and researchers with a valuable basis for identifying regional trends, underlying risk factors, and variations in prevalence. This information is crucial for tailoring prevention strategies, refining diagnostic approaches, and guiding future research efforts aimed at addressing the global burden of FSGS.

5. Declarations

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5.1. Author Contributions: Conceptualization, COC and VCV; Methodology, VCV; Formal analysis, COC, VCV and FALR. Investigation, COC and VCV; Data curation, VCV and FALR; Writing-original draft preparation, VCV, AGG and FALR; Writing-review and editing, COC, VCV, AGG and FALR; Project administration, VCV. All authors have read and agreed to the published version of the manuscript.

Data availability statement. Data will be made available on request.

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References

- 1. Suresh V, Stillman IE, Campbell KN, Meliambro K. Focal Segmental Glomerulosclerosis. Advances in Kidney Disease and Health. 2024 Jul;31(4):275–89.
- 2. Rosenberg AZ, Kopp JB. Focal Segmental Glomerulosclerosis. Clinical Journal of the American Society of Nephrology. 2017 Mar;12(3):502–17.
- 3. D'Agati VD, Fogo AB, Bruijn JA, Jennette JC. Pathologic classification of focal segmental glomerulosclerosis: a working proposal. American Journal of Kidney Diseases. 2004 Feb;43(2):368–82.
- 4. Bonilla M, Efe O, Selvaskandan H, Lerma E V., Wiegley N. A Review of Focal Segmental Glomerulosclerosis Classification With a Focus on Genetic Associations. Kidney Med. 2024 Jun;6(6):100826.
- 5. Hattori M. Anti-nephrin autoantibodies: novel predictors of post-transplant recurrence of focal segmental glomerular sclerosis. Kidney Int. 2024 Oct;106(4):570–2.
- 6. D'Agati VD, Kaskel FJ, Falk RJ. Focal Segmental Glomerulosclerosis. New England Journal of Medicine. 2011 Dec 22;365(25):2398–411.
- 7. Rood IM, Deegens JKJ, Wetzels JFM. Genetic causes of focal segmental glomerulosclerosis: implications for clinical practice. Nephrology Dialysis Transplantation. 2012 Mar 1;27(3):882–90.
- 8. Bai J, Yin X, Li J, Li J, Niu Y, Li Z, et al. Incidence, risk factors, and outcomes of recurrent focal segmental glomerulosclerosis in pediatric kidney transplant recipients: A systematic review and meta-analysis. Clin Transplant. 2023 Nov 19;37(11).
- 9. Ekrikpo UE, Obiagwu PN, Udo AI, Chukwuonye II, Noubiap JJ, Okpechi-Samuel US, et al. Prevalence and distribution of primary glomerular diseases in Africa: a systematic review and meta-analysis of observational studies. Pan African Medical Journal. 2023;45.
- 10. Ulasi II, Tzur S, Wasser WG, Shemer R, Kruzel E, Feigin E, et al. High Population Frequencies of APOL1 Risk Variants Are Associated with Increased Prevalence of Non-Diabetic Chronic Kidney Disease in the Igbo People from South-Eastern Nigeria. Nephron Clin Pract. 2013 Jul 13;123(1–2):123–8.
- 11. Zhao H, Ma L, Yan M, Wang Y, Zhao T, Zhang H, et al. Association between *MYH9* and *APOL1* Gene Polymorphisms and the Risk of Diabetic Kidney Disease in Patients with Type 2 Diabetes in a Chinese Han Population. J Diabetes Res. 2018;2018:1–6.
- 12. Ko GJ, Rhee CM, Kalantar-Zadeh K, Joshi S. The Effects of High-Protein Diets on Kidney Health and Longevity. Journal of the American Society of Nephrology. 2020 Aug;31(8):1667–79.
- de Boer IH, Zelnick LR, Ruzinski J, Friedenberg G, Duszlak J, Bubes VY, et al. Effect of Vitamin D and Omega-3 Fatty Acid Supplementation on Kidney Function in Patients With Type 2 Diabetes. JAMA. 2019 Nov 19;322(19):1899.

- 14. Bakis H, Chauveau P, Combe C, Pfirmann P. Mediterranean Diet for Cardiovascular Risk Reduction in Chronic Kidney Disease. Advances in Kidney Disease and Health. 2023 Nov;30(6):496–501.
- 15. Yau K, Kuah R, Cherney DZI, Lam TKT. Obesity and the kidney: mechanistic links and therapeutic advances. Nat Rev Endocrinol. 2024 Jun 13;20(6):321–35.
- 16. McGrogan A, Franssen CFM, de Vries CS. The incidence of primary glomerulonephritis worldwide: a systematic review of the literature. Nephrology Dialysis Transplantation. 2011 Feb 1;26(2):414–30.
- 17. Jegatheesan D, Nath K, Reyaldeen R, Sivasuthan G, John GT, Francis L, et al. Epidemiology of biopsy-proven glomerulonephritis in <scp>Q</scp> ueensland adults. Nephrology. 2016 Jan 23;21(1):28–34.
- 18. Zink CM, Ernst S, Riehl J, Helmchen U, Gröne HJ, Floege J, et al. Trends of renal diseases in Germany: review of a regional renal biopsy database from 1990 to 2013. Clin Kidney J. 2019 Dec 1;12(6):795–800.
- 19. Garau M, Cabrera J, Ottati G, Caorsi H, Gonzalez Martinez F, Acosta N, et al. Temporal trends in biopsy proven glomerular disease in Uruguay, 1990-2014. PLoS One. 2018 Oct 29;13(10):e0206637.
- 20. Schena FP. Survey of the Italian Registry of Renal Biopsies. Frequency of the renal diseases for 7 consecutive years. The Italian Group of Renal Immunopathology. Nephrology Dialysis Transplantation. 1997 Mar 1;12(3):418–26.
- 21. Molnár A, Thomas MJ, Fintha A, Kardos M, Dobi D, Tislér A, et al. Kidney biopsy-based epidemiologic analysis shows growing biopsy rate among the elderly. Sci Rep. 2021 Dec 29;11(1):24479.
- 22. Nakagawa N, Kimura T, Sakate R, Wada T, Furuichi K, Okada H, et al. Demographics and treatment of patients with primary nephrotic syndrome in Japan using a national registry of clinical personal records. Sci Rep. 2023 Sep 7;13(1):14771.
- 23. Arias LF, Henao J, Giraldo RD, Carvajal N, Rodelo J, Arbeláez M. Glomerular diseases in a Hispanic population: review of a regional renal biopsy database. Sao Paulo Medical Journal. 2009;127(3):140–4.
- 24. Santangelo L, Netti GS, Giordano P, Carbone V, Martino M, Torres DD, et al. Indications and results of renal biopsy in children: a 36-year experience. World Journal of Pediatrics. 2018 Apr 22;14(2):127–33.
- 25. Ali A, Ali MU, Akhtar SZ. Histological pattern of paediatric renal diseases in northern Pakistan. J Pak Med Assoc. 2011 Jul;61(7):653–8.
- 26. Sadaf A, Khemchand M, Fouzia L, Asia Z. Clinicopathological profile of pediatric renal biopsies at a tertiary care hospital, Pakistan. Saudi Journal of Kidney Diseases and Transplantation. 2018;29(6):1403.

- 27. Yadav S, Kandalkar B. Epidemiology of Pediatric Renal Diseases and its Histopathological Spectrum A Single-Center Experience from India. Saudi Journal of Kidney Diseases and Transplantation. 2021;32(6):1744.
- 28. Wine R, Vasilevska-Ristovska J, Banh T, Knott J, Noone D, Gbadegesin R, et al. Trends in the epidemiology of childhood nephrotic syndrome in Africa: A systematic review. Glob Epidemiol. 2021 Nov;3:100061.
- 29. Li LS, Liu ZH. Epidemiologic data of renal diseases from a single unit in China: Analysis based on 13,519 renal biopsies. Kidney Int. 2004 Sep;66(3):920–3.
- 30. Zhang X, Liu S, Tang L, Wu J, Chen P, Yin Z, et al. Analysis of pathological data of renal biopsy at one single center in China from 1987 to 2012. Chin Med J (Engl). 2014;127(9):1715–20.
- 31. Zhou F d., Zhao M h., Zou W z., Liu G, Wang H. The changing spectrum of primary glomerular diseases within 15 years: A survey of 3331 patients in a single Chinese centre. Nephrology Dialysis Transplantation. 2008 Oct 8;24(3):870–6.
- 32. Xu X, Ning Y, Shang W, Li M, Ku M, Li Q, et al. Analysis of 4931 renal biopsy data in central China from 1994 to 2014. Ren Fail. 2016 Aug 8;38(7):1021–30.
- 33. Zhou Q, Yang X, Wang M, Wang H, Zhao J, Bi Y, et al. Changes in the diagnosis of glomerular diseases in east China: a 15-year renal biopsy study. Ren Fail. 2018 Oct 15;40(1):657–64.
- 34. Li Y, Yu X, Zhang W, Lv J, Lan P, Wang Z, et al. Epidemiological characteristics and pathological changes of primary glomerular diseases. PLoS One. 2022 Aug 18;17(8):e0272237.
- 35. Hou JH, Zhu HX, Zhou ML, Le WB, Zeng CH, Liang SS, et al. Changes in the Spectrum of Kidney Diseases: An Analysis of 40,759 Biopsy-Proven Cases from 2003 to 2014 in China. Kidney Diseases. 2018;4(1):10–9.
- 36. Nie S, He W, Huang T, Liu D, Wang G, Geng J, et al. The Spectrum of Biopsy-Proven Glomerular Diseases among Children in China. Clinical Journal of the American Society of Nephrology. 2018 Jul;13(7):1047–54.
- 37. Su S, Yu J, Wang Y, Wang Y, Li J, Xu Z. Clinicopathologic correlations of renal biopsy findings from northeast China. Medicine. 2019 Jun;98(23):e15880.
- 38. Wang Y, Zhang L, Yuan L, Xie Q, Liu S, Hao C ming. Changes in the spectrum of biopsy-proven renal diseases over 11 years: a single-center study in China. Ren Fail. 2024 Dec 31;46(2).
- 39. Nie P, Chen R, Luo M, Dong C, Chen L, Liu J, et al. Clinical and Pathological Analysis of 4910 Patients Who Received Renal Biopsies at a Single Center in Northeast China. Biomed Res Int. 2019 Mar 26;2019:1–9.

- 40. Hu YC, Feng YX, Lv XA, Wang R. A Clinical and Pathological Analysis of 3722 Renal Biopsy Specimens from Adults with Primary Glomerular Disease in Shandong Province, China. WIMJ Open. 2014;1(2).
- 41. Hu R, Quan S, Wang Y, Zhou Y, Zhang Y, Liu L, et al. Spectrum of biopsy proven renal diseases in Central China: a 10-year retrospective study based on 34,630 cases. Sci Rep. 2020 Jul 3;10(1):10994.
- 42. Huang Y, Shi K, Zhu X, Yuan S, Chen X, Fu X, et al. Disease spectrum of 9 310 cases of renal biopsy pathological diagnosis from a single center in China. Zhong Nan Da Xue Xue Bao Yi Xue Ban. 2022 May 28;47(5):546–54.
- 43. Chen H, Tang Z, Zeng C, Hu W, Wang Q, Yu Y, et al. Pathological demography of native patients in a nephrology center in China. Chin Med J (Engl). 2003 Sep;116(9):1377–81.
- 44. Xiong M, Wang L, Liu X, Yue S, Dong J, Li Y, et al. Kidney Biopsies in Elderly Chinese Patients: A Nationwide Survey. American Journal of Kidney Diseases. 2020 Aug;76(2):295–7.
- 45. Dong J, Li Y, Yue S, Liu X, Wang L, Xiong M, et al. The profiles of biopsy-proven renal tubulointerstitial lesions in patients with glomerular disease. Ann Transl Med. 2020 Sep;8(17):1066–1066.
- 46. Das U, Dakshinamurty K, Prayaga A. Pattern of biopsy-proven renal disease in a single center of south India: 19 years experience. Indian J Nephrol. 2011;21(4):250.
- 47. Imtiaz S, Nasir K, Drohlia M, Salman B, Ahmad A. Frequency of kidney diseases and clinical indications of pediatric renal biopsy: A single center experience. Indian J Nephrol. 2016;26(3):199.
- 48. Mittal P, Agarwal SK, Singh G, Bhowmik D, Mahajan S, Dinda A, et al. Spectrum of biopsy-proven renal disease in northern India: A single-centre study. Nephrology. 2020 Jan 6;25(1):55–62.
- 49. Sharma M, Mazumder M, Mahanta P, Doley P, Pegu G, Alam S, et al. Histological patterns of renal diseases in children: A 12-year experience from a single Tertiary Care Center in North-East India. Saudi Journal of Kidney Diseases and Transplantation. 2021;32(2):364.
- 50. Kanodia K V., Vanikar A V., Nigam LK, Patel RD, Suthar KS, Gera DN, et al. Pediatric Renal Biopsies in India: A Single-Centre Experience of Six Years. Nephrourol Mon. 2015 Jun 28;7(4).
- 51. Gopaliah L, Sudakaran I, Nalumakkal S, Narayanan R, Vareed B. Spectrum of biopsy-proven renal diseases: A single center experience. Saudi Journal of Kidney Diseases and Transplantation. 2018;29(2):392.
- 52. Golay V, Trivedi M, Abraham A, Roychowdhary A, Pandey R. The spectrum of glomerular diseases in a single center: A clinicopathological correlation. Indian J Nephrol. 2013;23(3):168.

- 53. Zahir Z, Wani A, Jain M, Agrawal V, Jain S. Pediatric glomerular diseases in North India–Epidemiology and clinicopathologic correlation. Indian J Nephrol. 2023;33(1):28.
- 54. Bhawane A, Pasari AS, Tolani P, Balwani MR. Spectrum of Biopsy-proven Native Kidney Disease in Central India. Saudi Journal of Kidney Diseases and Transplantation. 2022;33(5):688–92.
- 55. Sugiyama H, Yokoyama H, Sato H, Saito T, Kohda Y, Nishi S, et al. Japan Renal Biopsy Registry: the first nationwide, web-based, and prospective registry system of renal biopsies in Japan. Clin Exp Nephrol. 2011 Aug 1;15(4):493–503.
- 56. Yokoyama H, Sugiyama H, Sato H, Taguchi T, Nagata M, Matsuo S, et al. Renal disease in the elderly and the very elderly Japanese: analysis of the Japan Renal Biopsy Registry (J-RBR). Clin Exp Nephrol. 2012 Dec 11;16(6):903–20.
- 57. Yokoyama H, Sugiyama H, Narita I, Saito T, Yamagata K, Nishio S, et al. Outcomes of primary nephrotic syndrome in elderly Japanese: retrospective analysis of the Japan Renal Biopsy Registry (J-RBR). Clin Exp Nephrol. 2015 Jun 18;19(3):496–505.
- 58. Ozeki T, Maruyama S, Imasawa T, Kawaguchi T, Kitamura H, Kadomura M, et al. Clinical manifestations of focal segmental glomerulosclerosis in Japan from the Japan Renal Biopsy Registry: age stratification and comparison with minimal change disease. Sci Rep. 2021 Jan 28;11(1):2602.
- 59. Goto K, Imaizumi T, Hamada R, Ishikura K, Kosugi T, Narita I, et al. Renal pathology in adult and paediatric population of Japan: review of the Japan renal biopsy registry database from 2007 to 2017. J Nephrol. 2023 Aug 19;36(8):2257–67.
- 60. Sugiyama H, Yokoyama H, Sato H, Saito T, Kohda Y, Nishi S, et al. Japan Renal Biopsy Registry and Japan Kidney Disease Registry: Committee Report for 2009 and 2010. Clin Exp Nephrol. 2013 Apr 6;17(2):155–73.
- 61. Imtiaz S, Nasir K, Drohlia M, Salman B, Ahmad A. Frequency of kidney diseases and clinical indications of pediatric renal biopsy: A single center experience. Indian J Nephrol. 2016;26(3):199.
- 62. Safdar RS, Mehar MF, Asghar A, Buzdar N. Focal Segmental Glomerulosclerosis in Paediatric Population of South Punjab Pakistan: A Tertiary Care Hospital Experience. Pak J Med Sci. 2021 Feb 3;37(2).
- 63. Choi IJ, Jeong HJ, Han DS, Lee JS, Choi KH, Kang SW, et al. An analysis of 4,514 cases of renal biopsy in Korea. Yonsei Med J. 2001;42(2):247.
- 64. Chang JH, Kim DK, Kim HW, Park SY, Yoo TH, Kim BS, et al. Changing prevalence of glomerular diseases in Korean adults: a review of 20 years of experience. Nephrology Dialysis Transplantation. 2009 Aug 1;24(8):2406–10.
- 65. Shin HS, Cho DH, Kang SK, Kim HJ, Kim SY, Yang JW, et al. Patterns of renal disease in South Korea: a 20-year review of a single-center renal biopsy database. Ren Fail. 2017 Jan 1;39(1):540–6.

- 66. Yim T, Kim SU, Park S, Lim JH, Jung HY, Cho JH, et al. Patterns in renal diseases diagnosed by kidney biopsy: A single-center experience. Kidney Res Clin Pract. 2020 Mar 31;39(1):60–9.
- 67. Farahani E, Nili F, Moghimian M, Jahanzad I, Minoo F sadat, Abdollahi A, et al. Analysis of the Prevalence and Trend in Biopsy-Proven Native Kidney Diseases in Iranian Population: A 12-year Survey from a Referral Center. Iran J Pathol. 2023 Apr 1;18(2):202–9.
- 68. Daneshpajouhnejad P, Behzadi E, Amoushahi S, Aghabozorgi A, Farmani A, Hosseini SM, et al. A six-year survey of the spectrum of renal disorders on native kidney biopsy results in Central Iran and a review of literature. Saudi Journal of Kidney Diseases and Transplantation. 2018;29(3):658.
- 69. Pakfetrat M, Malekmakan L, Torabinezhad S, Yousefi O, Naddaffard D. Review of Renal Biopsies, A Single Center Experience. Iran J Kidney Dis. 2020 Jan;14(1):12–9.
- 70. Nili F, Farahani E, Moghimian M, Jahanzad I, Minoo FS, Salarvand S, et al. Spectrum and Distribution of Biopsy-proven Kidney Diseases: A 12-year Survey of a Single Center in Iran. Saudi Journal of Kidney Diseases and Transplantation. 2023;34(4):346–54.
- 71. Alhozali HM, Ahmed RA, Albeirouti RB, Alotibi FA, Ghazi DK, Shikdar MA, et al. Histopathological and Clinical Findings of Biopsy-Proven Focal and Segmental Glomerulosclerosis: A Retrospective Study. Cureus. 2022 Mar 11;
- 72. Abdullah LS. Histopathological pattern of pediatric renal diseases: a study from a university hospital in western Saudi Arabia. Saudi J Kidney Dis Transpl. 2012 Mar;23(2):377–84.
- 73. Aslam N, Khawaja N, Nawaz Z, Mushtaq F, Mousa D, Rehman E, et al. Pattern of glomerular disease in the Saudi population: A single-center, five-year retrospective study. Saudi Journal of Kidney Diseases and Transplantation. 2013;24(6):1265.
- 74. Demircin G, Delibaş A, Bek K, Erdoğan Ö, Bülbül M, Baysun Ş, et al. A one-center experience with pediatric percutaneous renal biopsy and histopathology in Ankara, Turkey. Int Urol Nephrol. 2009 Dec 12;41(4):933–9.
- 75. Fidan K, Isik Gonul I, Büyükkaragöz B, Isiyel E, Arinsoy T, Soylemezoglu O. Changing trends in pediatric renal biopsies: analysis of pediatric renal biopsies in national nephrology registry data. Ren Fail. 2016 Sep 13;38(8):1228–33.
- 76. Turkmen A, Sumnu A, Cebeci E, Yazici H, Eren N, Seyahi N, et al. Epidemiological features of primary glomerular disease in Turkey: a multicenter study by the Turkish Society of Nephrology Glomerular Diseases Working Group. BMC Nephrol. 2020 Dec 14;21(1):481.
- 77. Almardini R, Albaramki J, Al-Saliata G, Farah M, AlRabadi K, Albderat J. Pediatric focal segmental glomerulosclerosis in Jordan: A tertiary hospital experience. Saudi Journal of Kidney Diseases and Transplantation. 2018;29(4):816.
- 78. Sheyyab A, Al-thnaibat M, Zghayer AA, Alsheyyab J, Hamed R. Common Glomerular Diseases in Adult Jordanians: A Single-Center Experience. Int J Nephrol. 2022 Jul 4;2022:1–5.

- 79. Subedi M, Bartaula B, Pant AshokR, Adhikari P, Sharma SanjibK. Pattern of glomerular disease and clinicopathological correlation: A single-center study from Eastern Nepal. Saudi Journal of Kidney Diseases and Transplantation. 2018;29(6):1410.
- 80. Garyal, Kafle RK. Hisopathological spectrum of glomerular disease in nepal: a seven-year retrospective study. Nepal Med Coll J. 2008 Jun;10(2):126–8.
- 81. Khatri B, Baral A, Maharjan S, Khatri B. Primary Focal Segmental Glomerulosclerosis among Patients with Glomerular Disease Undergoing Kidney Biopsy in a Tertiary Care Centre: A Descriptive Cross-sectional Study. Journal of Nepal Medical Association. 2023 Feb 1;61(258):163–6.
- 82. Karnib HH, Gharavi AG, Aftimos G, Mahfoud Z, Saad R, Gemayel E, et al. A 5-year survey of biopsy proven kidney diseases in Lebanon: significant variation in prevalence of primary glomerular diseases by age, population structure and consanguinity. Nephrology Dialysis Transplantation. 2010 Dec 1;25(12):3962–9.
- 83. Aoun M, Halabi C, Ammar W. Treatment of Glomerular Diseases in Lebanon. Saudi Journal of Kidney Diseases and Transplantation. 2021;32(4):1089.
- 84. Yahya TM, Pingle A, Boobes Y, Pingle S. Analysis of 490 kidney biopsies: data from the United Arab Emirates Renal Diseases Registry. J Nephrol. 1998;11(3):148–50.
- 85. Chiu HF, Chen H chun, Lu KC, Shu KH. Distribution of glomerular diseases in Taiwan: preliminary report of National Renal Biopsy Registry—publication on behalf of Taiwan Society of Nephrology. BMC Nephrol. 2018 Dec 10;19(1):6.
- 86. Parichatikanond P, Chawanasuntorapoj R, Shayakul C, Choensuchon B, Vasuvattakul S, Vareesangthip K, et al. An analysis of 3,555 cases of renal biopsy in Thailand. J Med Assoc Thai. 2006 Aug;89 Suppl 2:S106-11.
- 87. Alwahaibi N, Al-Khazimi O, Al-Riyami M. Histopathological Spectrum of Glomerular Diseases in Oman: A Five-year Study. Saudi Journal of Kidney Diseases and Transplantation. 2022;33(3):425–31.
- 88. Ali AA, Sharif DA, Almukhtar SE, Abd KH, Saleem ZSM, Hughson MD. Incidence of glomerulonephritis and non-diabetic end-stage renal disease in a developing middle-east region near armed conflict. BMC Nephrol. 2018 Dec 11;19(1):257.
- 89. Habib M, Badruddoza S. Pattern of glomerular diseases among adults in Rajshahi, the Northern Region of Bangladesh. Saudi Journal of Kidney Diseases and Transplantation. 2012;23(4):876.
- 90. Muthukuda C, Suriyakumara V, Sosai C, Samarathunga T, Laxman M, Marasinghe A. Clinicopathological spectrum of biopsy-proven renal diseases of patients at a single center in Sri Lanka: a cross sectional retrospective review. BMC Nephrol. 2023 Jun 22;24(1):181.
- 91. AlYousef A, AlSahow A, AlHelal B, Alqallaf A, Abdallah E, Abdellatif M, et al. Glomerulonephritis Histopathological Pattern Change. BMC Nephrol. 2020 Dec 18;21(1):186.

- 92. Woo KT, Chan CM, Lim C, Choo J, Chin YM, Teng EWL, et al. A Global Evolutionary Trend of the Frequency of Primary Glomerulonephritis over the Past Four Decades. Kidney Diseases. 2019;5(4):247–58.
- 93. Stratta P, Segoloni GP, Canavese C, Sandri L, Mazzucco G, Roccatello D, et al. Incidence of biopsy-proven primary glomerulonephritis in an Italian Province. American Journal of Kidney Diseases. 1996 May;27(5):631–9.
- 94. Panichi V, Pasquariello A, Innocenti M, Meola M, Mantuano E, Beati S, et al. The Pisa experience of renal biopsies, 1977-2005. J Nephrol. 2007;20(3):329–35.
- 95. Gesualdo L, Di Palma AM, Morrone LF, Strippoli GF, Schena FP. The Italian experience of the national registry of renal biopsies. Kidney Int. 2004 Sep;66(3):890–4.
- 96. Zaza G, Bernich P, Lupo A. Incidence of primary glomerulonephritis in a large North-Eastern Italian area: a 13-year renal biopsy study. Nephrology Dialysis Transplantation. 2013 Feb 1;28(2):367–72.
- 97. Covic A, Schiller A, Volovat C, Gluhovschi G, Gusbeth-Tatomir P, Petrica L, et al. Epidemiology of renal disease in Romania: a 10 year review of two regional renal biopsy databases. Nephrology Dialysis Transplantation. 2006 Feb 1;21(2):419–24.
- 98. Volovăt C, Căruntu I, Costin C, Stefan A, Popa R, Volovăt S, et al. Changes in the histological spectrum of glomerular diseases in the past 16 years in the North-Eastern region of Romania. BMC Nephrol. 2013 Dec 15;14(1):148.
- 99. Covic A, Vlad CE, Căruntu ID, Voroneanu L, Hogas S, Cusai S, et al. Epidemiology of biopsy-proven glomerulonephritis in the past 25 years in the North-Eastern area of Romania. Int Urol Nephrol. 2022 Feb 15;54(2):365–76.
- 100. Kurnatowska I, Jędrzejka D, Małyska A, Wągrowska-Danilewicz M, Danilewicz M, Nowicki M. Trends in the Incidence of Biopsy-Proven Glomerular Diseases in the Adult Population in Central Poland in the Years 1990-2010. Kidney Blood Press Res. 2012;35(4):254–8.
- 101. Perkowska-Ptasinska A, Bartczak A, Wagrowska-Danilewicz M, Halon A, Okon K, Wozniak A, et al. Clinicopathologic correlations of renal pathology in the adult population of Poland. Nephrology Dialysis Transplantation. 2017 Apr 1;32(suppl_2):ii209–18.
- 102. Rivera F. Frequency of renal pathology in Spain 1994-1999. Nephrology Dialysis Transplantation. 2002 Sep 1;17(9):1594–602.
- 103. López-Gómez JM, Rivera F. Registro de glomerulonefritis de la Sociedad Española de Nefrología en 2019: pasado, presente y nuevos retos. Nefrología. 2020 Jul;40(4):371–83.
- 104. Jönsson A, Hellmark T, Segelmark M, Forsberg A, Dreja K. Causes of nephrotic syndrome in Sweden: The relevance of clinical presentation and demographics. Frontiers in Nephrology. 2023 Mar 17;3.

- 105. Rychlik I, Jancova E, Tesar V, Kolsky A, Lacha J, Stejskal J, et al. The Czech registry of renal biopsies. Occurrence of renal diseases in the years 1994-2000. Nephrology Dialysis Transplantation. 2004 Dec 1;19(12):3040–9.
- 106. Maixnerova D, Jancova E, Skibova J, Rysava R, Rychlik I, Viklicky O, et al. Nationwide biopsy survey of renal diseases in the Czech Republic during the years 1994–2011. J Nephrol. 2015 Feb 23;28(1):39–49.
- 107. Heaf J, Løkkegaard H, Larsen S. The epidemiology and prognosis of glomerulonephritis in Denmark 1985–1997. Nephrology Dialysis Transplantation. 1999 Aug 1;14(8):1889–97.
- 108. Heaf JG, Sørensen SS, Hansen A. Increased incidence and improved prognosis of glomerulonephritis: a national 30-year study. Clin Kidney J. 2021 May 28;14(6):1594–602.
- 109. Naumovic R, Pavlovic S, Stojkovic D, Basta-Jovanovic G, Nesic V. Renal biopsy registry from a single centre in Serbia: 20 years of experience. Nephrology Dialysis Transplantation. 2008 Oct 8;24(3):877–85.
- 110. Paripovic D, Kostic M, Kruscic D, Spasojevic B, Lomic G, Markovic-Lipkovski J, et al. Indications and results of renal biopsy in children: a 10-year review from a single center in Serbia. J Nephrol. 2012;25(6):1054–9.
- 111. Horvatic I, Tisljar M, Bulimbasic S, Bozic B, Galesic Ljubanovic D, Galesic K. Epidemiologic data of adult native biopsy-proven renal diseases in Croatia. Int Urol Nephrol. 2013 Dec 1;45(6):1577–87.
- 112. Laurens W, Deleersnijder D, Dendooven A, Lerut E, De Vriese AS, Dejagere T, et al. Epidemiology of native kidney disease in Flanders: results from the FCGG kidney biopsy registry. Clin Kidney J. 2022 Jun 23;15(7):1361–72.
- 113. Deleersnijder D, Knops N, Trouet D, Van Hoeck K, Karamaria S, Vande Walle J, et al. Epidemiology and clinicopathological characteristics of native kidney disease in children in Flanders, Belgium. Pediatric Nephrology. 2023 May 13;38(5):1533–45.
- 114. Simon P, Ramée MP, Autuly V, Laruelle E, Charasse C, Cam G, et al. Epidemiology of primary glomerular diseases in a French region. Variations according to period and age. Kidney Int. 1994 Oct;46(4):1192–8.
- 115. Braun N, Schweisfurth A, Lohöfener C, Lange C, Gründemann C, Kundt G, et al. Epidemiology of glomerulonephritis in Northern Germany. Int Urol Nephrol. 2011 Dec 20;43(4):1117–26.
- 116. Oygar DD, Neild GH. Reporting renal biopsies from Cyprus: a systematic approach. J Nephropathol. 2017 Apr 13;6(3):231–9.
- 117. Brazdziute E, Miglinas M, Gruodyte E, Priluckiene J, Tamosaitis A, Bumblyte IA, et al. Nationwide renal biopsy data in Lithuania 1994–2012. Int Urol Nephrol. 2015 Apr 17;47(4):655–62.

- 118. Riispere Ž, Ots-Rosenberg M. Occurrence of kidney diseases and patterns of glomerular disease based on a 10-year kidney biopsy material: A retrospective single-centre analysis in Estonia. Scand J Urol Nephrol. 2012 Oct 1;46(5):389–94.
- 119. Sousa P, Brás C, Menezes C, Vizcaino R, Costa T, Faria MS, et al. Biópsias renais percutâneas em crianças: uma revisão de 24 anos em um centro terciário no norte de Portugal. Brazilian Journal of Nephrology. 2024 Sep;46(3).
- 120. Wirta O, Mustonen J, Helin H, Pasternack A. Incidence of biopsy-proven glomerulonephritis. Nephrology Dialysis Transplantation. 2007 Aug 17;23(1):193–200.
- 121. Gjerstad AC, Skrunes R, Tøndel C, Åsberg A, Leh S, Klingenberg C, et al. Kidney biopsy diagnosis in childhood in the Norwegian Kidney Biopsy Registry and the long-term risk of kidney replacement therapy: a 25-year follow-up. Pediatric Nephrology. 2023 Apr 22;38(4):1249–56.
- 122. Hanko JB, Mullan RN, O'Rourke DM, McNamee PT, Maxwell AP, Courtney AE. The changing pattern of adult primary glomerular disease. Nephrology Dialysis Transplantation. 2009 Oct;24(10):3050–4.
- 123. Swaminathan S, Leung N, Lager DJ, Melton LJ, Bergstralh EJ, Rohlinger A, et al. Changing Incidence of Glomerular Disease in Olmsted County, Minnesota. Clinical Journal of the American Society of Nephrology. 2006 May;1(3):483–7.
- 124. Hommos MS, De Vriese AS, Alexander MP, Sethi S, Vaughan L, Zand L, et al. The Incidence of Primary vs Secondary Focal Segmental Glomerulosclerosis: A Clinicopathologic Study. Mayo Clin Proc. 2017 Dec;92(12):1772–81.
- 125. Sim JJ, Batech M, Hever A, Harrison TN, Avelar T, Kanter MH, et al. Distribution of Biopsy-Proven Presumed Primary Glomerulonephropathies in 2000-2011 Among a Racially and Ethnically Diverse US Population. American Journal of Kidney Diseases. 2016 Oct;68(4):533–44.
- 126. Nair R, Walker PD. Is IgA nephropathy the commonest primary glomerulopathy among young adults in the USA? Kidney Int. 2006 Apr;69(8):1455–8.
- 127. Murugapandian S, Mansour I, Hudeeb M, Hamed K, Hammode E, Bijin B, et al. Epidemiology of Glomerular Disease in Southern Arizona. Medicine. 2016 May;95(18):e3633.
- 128. Machado SGR, Quadros T, Watanabe Y, Aquino CF, Otoni A, Pinto SW. Most common histopathological patterns of the Minas Gerais Association of the Centers of Nephrology. Rev Assoc Med Bras. 2019 Mar;65(3):441–5.
- 129. Polito MG, de Moura LAR, Kirsztajn GM. An overview on frequency of renal biopsy diagnosis in Brazil: clinical and pathological patterns based on 9617 native kidney biopsies. Nephrology Dialysis Transplantation. 2010 Feb 1;25(2):490–6.
- 130. Costa DM do N, Valente LM, Gouveia PA da C, Sarinho FW, Fernandes GV, Cavalcante MAG de M, et al. Comparative analysis of primary and secondary glomerulopathies in the

- northeast of Brazil: data from the Pernambuco Registry of Glomerulopathies REPEG. Jornal Brasileiro de Nefrologia. 2017;39(1).
- 131. Malafronte P, Mastroianni-Kirsztajn G, Betonico GN, Romao JE, Alves MAR, Carvalho MF, et al. Paulista registry of glomerulonephritis: 5-year data report. Nephrology Dialysis Transplantation. 2006 Sep 12;21(11):3098–105.
- 132. Thomé GG, Bianchini T, Bringhenti RN, Schaefer PG, Barros EJG, Veronese FV. The spectrum of biopsy-proven glomerular diseases in a tertiary Hospital in Southern Brazil. BMC Nephrol. 2021 Dec 13;22(1):414.
- 133. Thomé GG, Bianchini T, Bringhenti RN, Schaefer PG, Barros EJG, Veronese FV. The spectrum of biopsy-proven glomerular diseases in a tertiary Hospital in Southern Brazil. BMC Nephrol. 2021 Dec 13;22(1):414.
- 134. Arias LF, Jiménez CA, Arroyave MJ. Histologic variants of primary focal segmental glomerulosclerosis: presentation and outcome. Jornal Brasileiro de Nefrologia. 2013;35(2):112–9.
- 135. Barrera-Herrera LE, López Panqueva R del P, Flórez Vargas AA, Andrade Pérez RE. The spectrum of glomerular disease between the years 2003 and 2015 in Columbia: A review of 12,613 cases. Revista Española de Patología. 2017 Jan;50(1):3–7.
- 136. Prada Rico M, Rodríguez Cuellar CI, Fernandez Hernandez M, González Chaparro LS, Prado Agredo OL, Gastelbondo Amaya R. Characterization and Etiopathogenic Approach of Pediatric Renal Biopsy Patients in a Colombian Medical Center from 2007-2017. Int J Nephrol. 2018 Jun 28;2018:1–9.
- 137. Aroca-Martínez G, González-Torres HJ, Domínguez-Vargas A, García-Tolosa R, Castillo-Parodi L, Conde-Manotas J, et al. Glomerular Diseases in the Colombian Caribbean. Saudi Journal of Kidney Diseases and Transplantation. 2022 Feb;33(Suppl 1):S18–29.
- 138. Chávez Valencia V, Orizaga de La Cruz C, Becerra Fuentes JG, Fuentes Ramírez F, Parra Michel R, Aragaki Y, et al. [Epidemiology of glomerular disease in adults: a database review]. Gac Med Mex. 2014;150(5):403–8.
- 139. Cunningham A, Benediktsson H, Muruve DA, Hildebrand AM, Ravani P. Trends in Biopsy-Based Diagnosis of Kidney Disease: A Population Study. Can J Kidney Health Dis. 2018 Jan 20;5.
- 140. Valjalo R, Mallea MT. Caracterización de enfermedades glomerulares: análisis de 22 años de biopsias renales. Rev Med Chil. 2023 Feb;151(1):52–60.
- 141. Mazzuchi N, Acosta N, Caorsi H, Schwedt E, Di Martino LA, Mautone M, et al. [Frequency of diagnosis and clinic presentation of glomerulopathies in Uruguay]. Nefrologia. 2005;25(2):113–20.
- 142. Hurtado A, Escudero E, Stromquist CS, Urcia J, Hurtado ME, Gretch D, et al. Distinct patterns of glomerular disease in Lima, Peru. Clin Nephrol. 2000 May;53(5):325–32.

- 143. Asinobi AO, Ademola AD, Okolo CA, Yaria JO. Trends in the histopathology of childhood nephrotic syndrome in Ibadan Nigeria: preponderance of idiopathic focal segmental glomerulosclerosis. BMC Nephrol. 2015 Dec 15;16(1):213.
- 144. Onwubuya I, Adelusola K, Sabageh D, Ezike K, Olaofe O. Biopsy-proven renal disease in Ile-Ife, Nigeria: A histopathologic review. Indian J Nephrol. 2016;26(1):16.
- 145. Ibrahim S, Fadda S, Fayed A, Belal D. A five-year analysis of the incidence of glomerulonephritis at Cairo University Hospital-Egypt. Saudi Journal of Kidney Diseases and Transplantation. 2012;23(4):866.
- 146. Okpechi I, Swanepoel C, Duffield M, Mahala B, Wearne N, Alagbe S, et al. Patterns of renal disease in Cape Town South Africa: a 10-year review of a single-centre renal biopsy database. Nephrology Dialysis Transplantation. 2011 Jun 1;26(6):1853–61.
- 147. Aatif T, Maoujoud O, Montasser D, Benyahia M, Oualim Z. Glomerular diseases in the Military Hospital of Morocco: Review of a single centre renal biopsy database on adults. Indian J Nephrol. 2012;22(4):257.
- 148. Abdelraheem MB, Ali ETMA, Mohamed RM, Hassan EG, Abdalla OA, Mekki SO, et al. Pattern of glomerular diseases in Sudanese children: a clinico-pathological study. Saudi J Kidney Dis Transpl. 2010 Jul;21(4):778–83.
- 149. Hoy WE, Samuel T, Mott SA, Kincaid-Smith PS, Fogo AB, Dowling JP, et al. Renal biopsy findings among Indigenous Australians: a nationwide review. Kidney Int. 2012 Dec;82(12):1321–31.
- 150. Painter D, Clouston D, Ahn E, Kirwan P, Ledoux F, Tivollier JM, et al. The pattern of glomerular disease in new caledonia: preliminary findings. Pathology. 1996;28(1):32–5.

Figure Legends

Figure 1. PRISMA flowchart depicting the study selection process.

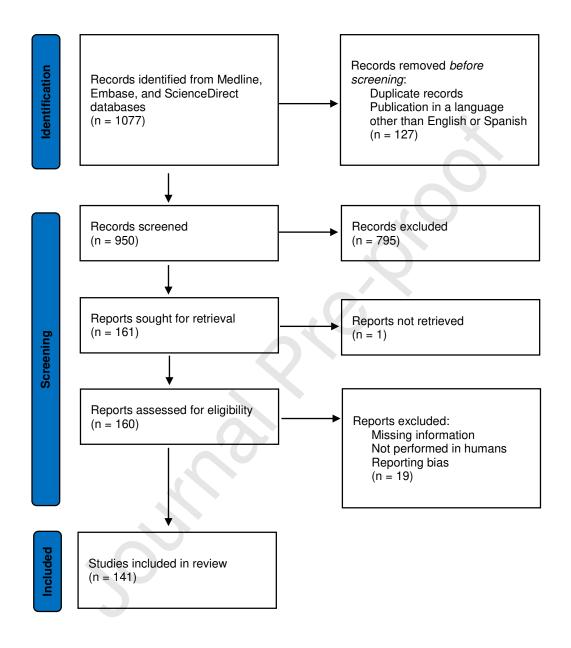


Figure 2. Prevalence of FSGS in the studied countries. A blue color scale represents prevalence, with darker shades indicating higher rates. Countries shown in grey lack available prevalence data.



Figure 3. Trends in prevalence of FSGS over time in selected countries.

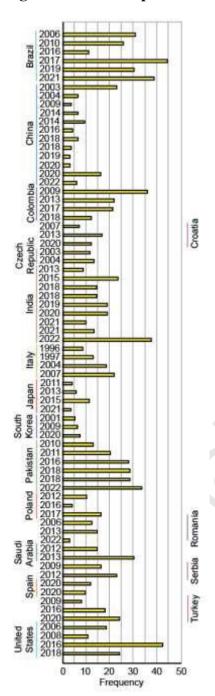


Table 1. Summary of the studies analyzed in this paper to estimate the worldwide prevalence of focal segmental glomerulosclerosis (FSGS).

Continent	Country	Period and reference	No. Biopsies	Prevalence (%)	Population	Description
		1979- 2002 ²⁹	13519	6	Adults. A single unit in China. and 57.3% were male.	Average age was 32.7 ± 12.2 (9 to 83) years. Biopsies with PGD 9278, FSGS was present in 557 cases
Asia	China	1987- 2012 ³⁰ 1993- 2017 ³¹	11618	5.82	General; All patients were Asian, 6 646 male cases and 4 972 female cases (1.33:1).	PGD 8,209 cases, median age of 35 years (range 3-85 years old). Among the PGD FSGS was present in 458 patients of 8209.
4			5398	Prevalence of FSGS was 2.7 in 1st period (1993- 1997), 3.3 in 2nd (1998-2002) and 3.8 in 3rd period (2003-2007).	Adults	One center, patients (age ≥14 years) were included, 3331 PGD, 110 FSGS (71 male/39 female).
		1994- 2014 ³²	4931	The proportion of FSGS increased from 1.74% in period 1(1994-1999) and 1.55% in period 2 (2000-	Adults	Ten hospitals in China. Average age was 35.2 years, 2254 patients (45.7%) were men. PGD

	1		2004) 4 5 5767 :		01.550
			2004) to 5.57% in		were 81.55%,
			period 3 (2005-		FSGS was 146
			2009), but then it		patients (3.63%),
			decreased to 3.09%		85 male and 61
			in period 4 (201-		female.
			2014).		
					A single center,
	2001-				patients aged (≥15
		10779	2.5	General	years old. Average
	2015 ³³				age was 40 ± 14.8
					years old
			FSGS was in 477		
			patients (5.05%),		
			group 1		
			(01/01/2001 to		
			12/21/2004) was		
			51 patients		
			(5.06%), group 2		Average age was
			(01/01/2005 to		41.2 years for
			12/31/2009) was		males and 41.60
	2001-	9448	109 patients	Adults	years or females.
	2019^{34}	7110	(6.04%), group 3	Tadito	60.39% male.
			(0.01/0), group 3 (01/01/2010 to		00.39 % maie.
			12/31/2014) was		
			205 patients		
			(6.26%), and group		
			4 (01/01/2015 to		
			12/31/2019) was		
			112 patients		
			(3.33%).		
			FSGS was 6% in		Patients > 14
			period 1979-2002		years. Renal
	2003- 2014 ³⁵	40759	and changes	Adults	Biopsy Registry of
			frequencies at		the National
	2011		7.34% to period of		Clinical Research
			7.54 % to period of		Center of Kidney
			2003-2014.		center of Ridney

Г	1	1			26 50 1
					age was 36.59 ±
					14.12 years. 52%
					male. PGD was
					67.1% of cases.
			FSGS was 6% in		
			male and 4% in		
			female patients.		
			5% in patients to 0-		
			12 year and 6% in	C.	
			13-18 year. FSGS		
			showed a		
			significant		Average age was
	2004-		decreasing trend,		13.5±4.1 years, 64% boys. Included 115 hospitals.
	2014 36	7962	starting at 14% in	Children	
			the first period		
		<	(2004-2007) and		
			decreasing to 6%		
			and 4% in the		
			second (2008-		
			2011) and third		
			periods (2012-		
			2014).		
					One hospital in
					northeast China.
	2007-				Only >14 years.
	2016 37	2725	2.68	Adults	Average age was
	2010				41.2±15.1 years.
					FSGS was 73
					patients.
					A single center,
					retrospective study
	2008-	10000			of native kidney
	2018 38	10996	8	General	biopsies, PGD was
	2008-				69.42%, with 8%
					for GSFS.
					3,593 cases of
	2017 39	4910	3.1	General	PGD (73.2%).
	2017				1 00 (13.270).

Т	1					A
						Average age was
						$42.6 \pm 15.7 \text{ years}$
						(range: 7 to 84
						years), 2629 males
						and 2281 females
						(ratio: 1.15:1).
						Shandong
						Province of China.
		2008-				FSGS patients
		2008 ⁻ 2013 ⁴⁰	3722	8.78	Adults	average was 37.9
		2013				± 12.5 years FSGS
						frequency was 327
						patients.
				FSGS frequency		
				was in 850 patients		
				(2.45%). FSGS had		Adults 54.27%
		2009-	35783	distribution peaks		male patients, ratio
				in the 20–39-year		male. Female
				age group. In	General	1.19:1, 31,256
		2018 41		elderly patients		were adults, with
				FSGS was 2.92%		age of adults
				and in children (-		40.8±15.2 years.
			14 years) FSGS			
				was 3.32%.		
						One center, two
						periods 2011-2015
						and 2016-2020.
						PGD were in
						66.93% cases
						(6166 cases),
		2011-	9310	14.97	Adults	among these
		2020 42	7510	- ··//	1100110	FSGS was 14.97%
						patients. In
						patients with age
						14-24 years FSGS
						was 14.04%, in
						25-44 years

Т		1		I	12.20%
					13.39% and 45-59
					years 18.33%.
					Included 17
					centers, of PGD
		1445			FSGS was 21.6%
					cases (133).
	2014-		21.6		Average age was
	2016 43			Adults	50.4±17.7 years,
	2010				55.6% male,
					14.3% had
					diabetes and
					50.4% had
					hypertension.
					Nationwide cross-
					sectional survey of
	2014-	7017	4.3	Adults	kidney biopsies in
	2018 44	7917	4.3		China. 59.6%
					male. FSGS was
					in 331 patients.
			FSGS was in 2,526 (4.4%) of patients, 199 (4.7%) pediatrics y 2,327 (4.4%) in adult patients	General	Retrospective
					study, including
		62569			56880 native
	2015-				biopsies at 1211
	2013-				hospitals across
	2017				China. Children
					were 4274 patients
					y adults 52606
					patients.
					Average age was
)				32.2 ± 18.3 years.
					1091 males and
	1990-				758 females.
	India 2008 46	1849	15.3	General	FSFGS was 195 of
	2008 10				1278 patients,
					median age 25
					years. Male 2.25:
					1 Female
	I	1		1	1

	I			I	A • 1 .
					A single center.
		423			Average age was
	1997-				$10.4 \pm 4.5 \text{ years},$
	2013 47		25.8	Children	57.9% male. PGD
	2013				was 360 patients,
					and of these FSGS
					was 109 patients.
			FOCO 41		A single center
	2006		FSGS was the	C.	study in northern
	2006-	3257	most common	Adults	India. Average age
	2016 48		PGD accounting		was 33.2 ± 14.2
			for 18.2%.		years, 61.9% male.
	2007				Median age was
	2007-	254	12.2	Children	15 years, 57%
	2018 49				female.
					Average age was
		335	8.02	Children	7.91 ± 3 years,
	2008- 2013 ⁵⁰				68.1% males.
					PGD was the most
					common in 81.8%,
					and FSGS was
					8.02%.
					Retrospective
		270	12.7	Adults	study. Average
					age was 36.92
	2009-				years. FSGS was
	2016 51	270	13.7		37 out of 270)
					most common
					PGD.
			22.58	Adults Children	Average age was
	2010-	409			$28 \pm 14.6 \text{ years.}$
	2012 52				Of 527 cases,
					primary FSGS was
					119 patients.
	2016- 2021 ⁵³				353 PGD, FSGS
					was present in 71
					patients.

						Mean age was
		2017- 2020 ⁵⁴	347			41.4 ± 15.7 years,
				4= 0-		58.5% were males.
				17.02	Adults	NS 36.3% of
						cases; 69% was
						PGD, FSGS was
						32 patients.
						Japan Renal
						Biopsy Registry.
						Prospective
						registry system, J-
		2007-	2126	3.62	Adults	RBR, 23 renal
		2008 55	2120	3.02	Adults	centers. Average
						age was 44.4 ± 21
						years, 1281 male.
						FSGS was present
						in 77 patients.
	Japan	2007- 2010 ⁵⁶	438	10.3	Adults	J-RBR. Mean age
						73 years, 226
						males. All with
						elderly (aged ≥65
						years) Japanese
						primary nephrotic
						syndrome (NS),
						reported FSGS in
						45 patients.
						J-RBR of the
						Japanese Society
				FSGS present in 99		of Nephrology
)			patients from 1259		included 1596
				patients in elderly		males. Renal
		2007-	2802	(≥65 years old) and	Adults	disease in the
		2011 57		83 patients (1.7%)		elderly (age ≥65
				in control group		years old) and
				(20-64 years) of 5021 cases.		very elderly (age
						≥80 years old)
						Japanese.
						Japanese.

		Ī				mpr :
						JRBR. Average
						age was 47 years,
					Adults	59.4% were male.
						The annual
						incidence of FSGS
		2007-	1409	3		accounted for 3.5-
		2016 58	1105	J	ridaits	4.5% Percentage
						of primary FSGS
						were constant at
						approximately 3%
						of all during 2007
						to 2016.
						Japan Renal
						Biopsy Registry
	2007-		32254			(J-RBR) 2007-
		2007-		3.4 in adults and	General	2017. 973 patient
		2017 59		3.7 in pediatrics	General	adults with FSGS
						and 129 pediatrics
						patients with
				Ť		FSGS.
						JRBR.
						Predominantly
						males 53.6% in
						2009 and 54.9% in
		1				2010, included
						4016 biopsies in
						2009 and 4681
		2009-	0.607	<i>5.</i> 2	4.1.1.	cases in 2010,
		2010 60	8697	5.3	Adults	average age
						(years) were
						46.7±19.9 and
						46.7±20.6
						respectively.
						From 7034, FSGS
						was present in 378
						patients.
						*

		l			<u> </u>	INICCC
						JNSCS patients
		2015				with primary NS;
		2015- 2018 ²²	6036	10.85	Adults	FSGS was 655
		2018				patients, age 39
						years, 57.5%
						males.
						Average age was
		1997-				10.48 ± 4.58
		2013 61	423	25.8	Children	years, 57.9% were
						males (245). FSGS
						in 109 patients.
						Pediatric kidney
		1998-		Primary FSGS in		biopsies between
		2005 25	415	18.3% of patients.	Children	3-15 years, male:
		2003		18.5% of patients.		female ratio was
						1.6:1.
				¹ 0		A tertiary care
						hospital. Average
						age in FSGS
	Pakistan	2010-	100	108 25.9	CI 'I I	patients was 7.0 ±
		2015 26	108		Children	4.2 years, 56.5%
						were male, FSGS
						was present in 28
						patients.
						. FSGS cases,
						mean age was
						8.8±3.0 years
				Primary FSGS in		while most of the
		2011-	307	40.4% (124	Children	children, 70
		2020 62		patients)		(56.5%) were
				• /		above 10 years of
						age; 64 cases were
						male.
				FSGS was present		
	South Korea	1973- 1995 ⁶³ 2097		in 154, Adults 97		82.5% PGD.
			(4.6%) and 57	General	Male: female ratio	
	Roica	1775		(4.0%) and 37 Children (4%).		was 1.5:1.
				Ciliureli (4 /0).		

	1987- 2006 ⁶⁴	1818	5.6	Adults	Average age was 36 years, male: female ratio was 1.02:1.
	1992- 2011 ⁶⁵	818 patients, adult group (18–59 years) included 758 cases, and the older group (≥60 years) included 60 cases.	FSGS was 3.5% in 18-59 years (N=621) and 12.8% in ≥60 years (N=43 patients)	Adults	Average age was 37.2 years, maleto-female ratio was 1.2:1;
	2001- 2013 ⁶⁶	1924	FSGS was in 130 patients (6.8%). People over 65 years of age, FSGS was in 18 patients out of 155 (11.6%).	Adults	A single center. Average age was 37.7 ± 16.5 years, 56% were male.
	2006- 2018 ⁶⁷	2975	15.9	Adults	The mean age of the patients was 27.4 years old; 51.6% were male.
Iran	2009- 2014 ⁶⁸	1054	24.8	Adults	Average age was 33.1 (±18.5) years, 43.3% were female.
	2011- 2017 ⁶⁹	774	188 patients were FSGS of 774 biopsies of primary	Adults	A Single Center. Average age was 33.9±17.5 years, 58% men.

			glomerulonephritis (24.2%).		
	2007- 2018 ⁷⁰	2975	15.9%	Adults	A Single Center. 51.6% was male. The mean age was 27.4 years.
Saudi	1989- 2020 ⁷¹	350	60 patients of 350 had FSGS (17.1%).	Children	A Retrospective Study. Included 39 pediatric patients and 21 adults. Male were 55% in both groups. The mean age of the pediatric patients was 7.13 ± 5.18 years, while that of the adult patients was 35.8 ± 14.3 years.
Arabia	1995- 2008 ⁷²	242	FSGS was 32 cases (13.2%)	Children	Only 183 patients had glomerular disease. Patients were 2 days to 17 years old, 122 were males.
30)	2005- 2009 ⁷³	348	27.6	General	Retrospective study, a single center, 176 were on adult males, 127 were on adult females and 45 were on children.
Turkey	1990- 2006 ⁷⁴	614	7.3	Children	376 with PGD (61.2%). Mean age was 10.4 years, 604 boys.

	1	ı	T		ī	
						Turkish Society of
						Nephrology
						Registry System
						(TSNRS), 2438
		1991-				were glomerular
		2010 75	3892	16.46	Children	diseases The age
		2010				range was 0 to 15
						years.
					C.	Male/female ratio
						was 1:1, FSGS
						was 641 patients.
						A multicenter
						study by the
						Turkish Society of
						Nephrology
				, (C)		Glomerular
						Diseases Working
						Group, 47 centers
		2000				of PGD of Turkish
		2009-	3875	21.9	Adults	Society of
		2019 76				Nephrology
						Glomerular
						Diseases (TSN-
						GOLD) Working
						Group. Average
						age was 41.5 ±
						14.9 years, 56.3%
						were male.
						Retrospective
						study of FSGS, a
						tertiary care
	Jordan	2010-	99	23	Children	hospital. Average
		2016 77				age 3.71± 2.59
						years, 66% were
						male.
		2013-				A Single-Center.
		2020 78	106	11.32	Adults	Average age was

						34 ±12.7 years,
						females were
						53.7%
						A single center.
						Tertiary care
						center. Average
						age was 35.3 ±
		2011		18.28% (18		13.5 years. 59.4%
		2014-	175	patients) had	Adults	female, ratio
		2016 ⁷⁹		FSGS.		female: male ratio
						of 1.5. The mean
						age of patients
				<		with FSGS was
Nepal	Nepai					36.89 years.
						Average age was
		2001- 2007 ⁸⁰				30.6 years for
						males and 32.9
			137	8	General	years for females.
						The male to
						female ratio was
						1.6: 1
		2022-	213	Primary FSGS in	Adults	One Center
		2022 81		10.33	ridaits	one center
						Average age was
				In Age ≤15 FSGS		$36.76 \pm 20 \text{ years}$
				was 6.6%, in 15<		(range 1–84),
		2003-		Age ≤60 FSGA		54.4% were male,
4		2007 82	1048	was 13.9% and age	General	20% pediatric
		·		>60 FSGS was		population, 67.3%
	Lebanon			12.1%.		young adult group
						and 12.7% elderly
						group.
		2014- 2015 ⁸³ 144		Children were		35 children (mean
			144	FSGS (5.7%) 31		age 11± 5.6 years)
				adult patients were	General	and 109 adults
			FSGS (28.4%)		(mean age 41.6	
				, ,		±16.5 years).

					74.2% were male
					in children group
					and 45.8% were
					male in adult
					patients,
					Data from the
					United Arab
XX 1. 1					Emirates Renal
United	1978-	400	10.2		Diseases Registry.
Arab	1996 ⁸⁴	490	18.3	Adults	378 PGD. Adults
Emirates					aged 14-66 years.
					FSGS were 69
					patients.
					National Renal
					Biopsy Registry
					with 17 medical
					centers. Average
	2014-		FSGS were in 133		age was 48.4 ±
Taiwan	2014-	1445	of 599 PGD	Adults	16.6 years, 53.8%
	2010		patients (25%),		cases male
					samples. Average
					age 50.4 ±17.1
					years, 55.6% male
					in FSGS group.
	1983-				A single center,
Thailand	2005 86	3355	2.8	Adults	FSGS was 61
	2003				patients.
					Pediatric kidney
	2011-		Primary FSGS in		biopsies between
Oman	2011-	596	18.3% patients.	Children	(3-15 years), male:
	2013		16.5% patients.		female ratio was
					1.6:1,
					Average age was
Iraq	2012-	662	22	Adults	27.3 ±17.6, 53%
maq	2013 88	002	22	Munts	were male. FSGS
					was 135 patients.
		•			

	Т					
	Panala dash	2008-	95	11.58	Adults	Average age was 30.29 years, 60% female; male to
	Bangladesh	2009 89	93	11.56	Adults	
						female ratio of
						1:1.5.
						One center, 55.7%
		2018-				females. Mean age
	Sri Lanka	2019 ⁹⁰	140	22.1	Adults	was 46±15.3
						years. FSGS in 31
						patients.
						Report of four
						public hospitals,
						356 adult native
						kidney biopsies,
						average age was
						39.8 (12 to 90
						years) years,
				10		61.2% were male.
				21.9		FSGS came third
						place with 78
						(21.9%) of the
						cases, 31 patients
	Kuwait	2013-	356		General	were less than 18
	Kuwaii	201891	330		General	years. 37 with sub-
						nephrotic
						proteinuria plus
						AKI, 15 with
						nephrotic
						syndrome, 11 with
						nephrotic
						syndrome plus
						AKI, 11 with sub-
						nephrotic
						proteinuria and 4
						with unexplained
						renal impairment.

	Singapore	1978- 2008 ⁹²	3,282 biopsies with PGD	11.88	General	Average age was 47.9 ± 13.5 years with a range from 15 to 85 years, predominantly in males in the first 3 decades. Retrospective study over 40 years. 390 cases of GSFS (11.88%). Frequency of FSGS was 5% in the 1st decade (1978-1988), 6% in 2nd decade (1988-1998), 15% in 3rd decade (1998-2008) and 25% in 4th decade (2008-2018).
Europe	Italy	1970- 1994 ⁹³	1926	FSGS present in 150 patients (7.8%), changes over time: 5.2% during 1907-1974, 6.3% 1975-1979, 6.7% 1980-1984, 9% in period 1985-1989 and 8.8% in period 1990-1994.	Adults	Mean age of patients undergoing biopsy (from 29.3± 12.2 years to 47.0 ± 17.8 years). In PGD predominance of males (> 2:1).
		1977- 2005 ⁹⁴	3269	19.8	Adults	66% PGD. Mean age 42 year, 59% males
		1979- 2014 ²⁴	213	11.6	General	A single tertiary pediatric hospital. Median 10.4 years

		1987- 1993 ²⁰	13835	FSGS was 11.8% of glomerulonephritis. The annual frequency of PGD to FSGS was 10.4% in 1993,	Adults	(range 0.6-24 years), 43.2% female. The Italian Group of Renal Immunopathology. Male sex was predominant in PGD (65%).
		1996- 2000 ⁹⁵	14607	FSGS was 16.9% of PGD with NS	Adults	Italian Immunopathology Group, Date from 128 rental units in Italy were reported. PGD were in 6,990 patients and were more frequent in males (64%),
		1998- 2010 ⁹⁶	4378	13.5	Adults	
	Romania	1995- 2004 ⁹⁷	635	11.5	General	Two regional renal biopsy databases. PGD was 401 cases. FSGS incidence was 10 p.m.p/year
Romania	Komama	2005- 2010 ⁹⁸	514	Overall, 288 patients, incidence of FSGS was 13.5 %, (0.51) p.m.p/year	Adults	Average age was 41.9±2.8 years, 58.5% were male. FSGS had an incidence of 0.70 p.m.p./year.

	I					
						Biopsy reports
						were divided into
						3 periods, 320
						from 1994 to 2004
						(period 1), 239
						from 2005 to 2010
						(period 2) and 442
				4.8% of PGD		biopsies between
		2011-	1101		Adults	2011 and 2019
		2019 99	1101	(2.9% of 442	Adults	(period 3). Mean
				cases)		age of the renal
					biopsy population	
						during period 3
						was 39.2 ±
						13.8 years, 65.2%
				, (C)		were male. PGD
						was 59.5% of the
						cases.
						Adults (>18
				•		years), in a single
			746	FSGS was 58 of 607 PGD patients	Adults	tertiary
						nephrology center
		1990-				serving an area of
		2010 100				Central Poland.
				(9.5%).		607 PGD. Average
						age was 40.5 ±8
						20.8 years, 411
	Poland					male patients.
						Polish Society of
						Nephrology. A
						total of 1939
						(21%) biopsies
	2009-	8843	15	General	were performed in	
	2014 101	0043			patients <18 years	
						of age, 6394
						(68.7%) in those
						18–64 and 955
						10 0. 4114 700

						(10.3%) in elderly
						individuals
						(defined as ≥65
						years of age).
						FSGS was present
						en 997 patients.
						Average age was
						47 years (19-87
						years) in FSGS
						patients and 55.1%
						were males.
						93 medical renal
						units. Median
				In children FSGS		(range) <15 years
				was 15.2% and 10.8% in adult	General	(487), 15–65 years
		1994- 1999 ¹⁰² 7				(4827) and >65
			7016	patients; in the		years (1510).
		1999		elderly FSGS was		Male/female ratio
	Spain			6%.		in children of 1.2,
				070.		in adults of 1.5
						and in the elderly
						of 1.4.
						The age range was
		1994-				15 to 65 years.
		2019 103	18852	9	Adults	Male/female ratio
						was 1.5.
		2014-				FSGS was present
	Sweden	2029 104	913	8.4	Adults	in 77 patients.
						Czech Registry of
			3294			Renal Biopsies
			biopsies in			included 28
	Czech Republic	1994-	adults and			centers. Mean age
		2000 105	710	10.8	General	10 years for
			biopsies in			children and 42
			< 15 years			years for adults,
						57.9% males.

		T			T	
						31 centers, 57.8%
						male. Mean age
						for children was
						10 years and for
						adults 44.5 years
		1994-	10472	12.6	General	FSGS incidence
		2011 106	10172	12.0	Concrai	was 3.9
						p.m.p/year, mean
						age for FSGS
						group was 40
						years, 56.8%
						males.
						Danish Renal
						Biopsy Register
						(DANYBIR).
					Adults	Average age was
		1985- 1997 ¹⁰⁷		13.66		42.6 ± 20.2 years.
			2380			FSGS was 325
						patients. Incidence
						5.7 pmp/year, age
	Denmark					FSGS group was
						43.4±19 and 34%
						were female.
						Danish Renal
						Biopsy Registry
		1985-	5042	7.02	A 1 14	and Patobank
		2014 108	5043	7.93	Adults	registries. FSGS
						was 400 patients
						of 5043 biopsies.
				FSGS was 15.1%		
		1007		causes of nephrotic		Average age was
		1987- 2006 ¹⁰⁹	1626	syndrome among	Adults	$39.1 \pm 13.8 \text{ years},$
	Serbia	2006 107		872 native kidney		51.2% were male.
				diseases.		
		2001				A single center.
		2001- 2010 ¹¹⁰	150	20.9	General	Mean age was
		2010 110				11.5 years, 56%
		<u> </u>			<u> </u>	

Departments, PGD was S7.4%.							were female
Croatia 1996- 2012 111 922 15.8 Adults Adults 16-84 years (range 16-84 years), patients were ≥16 years old. Median age 61.10 years, 62.1% males. 26 nephrology centers in Flanders (Belgium) 1152 glomerular final clinical diagnoses, FSGS incidence rate 12.1 p.m.p/year Diagnosis was often determined by results of genetic analysis Western France in the north of Brittany (Cotes d'Armor Depart ment)							patients. PGD was
Croatia 1996- 2012 11 922 15.8							_
Croatia 1996- 2012 11 922 15.8							Average age was
Croatia 1996- 2012 11 922 15.8 Adults 16-84 years), patients were ≥16 years old.							
2012 111		Croatia		922	15.8	Adults	
2017- 2019 112 2054 9.3 Adults Gelgium 1152 2017- 2019 112 2054 9.3 Adults Gelgium 1152 2017- 2020 13 148 11.1% Children Diagnosis was often determined by results of genetic analysis France 1976- 1990 114 480 10.6 General General General FSGS accounted 6.1% of all diagnoses (N=1,208). Over time FSGS was 4% in first period General children (≤15 years) and 1,185 Germany 1990- 2013 18 1208 time FSGS was 4% in first period General children (≤15 years) and 1,185 Germany 1990- 2013 18 1208 time FSGS was 4% General children (≤15 years) and 1,185 Germany 1,185 1,185 1,185 1,285			2012 111				-
Diagnosis was often determined by results of genetic analysis							_
2017- 2019 112 2054 9.3 Adults Gelgium 1152 glomerular final clinical diagnoses, FSGS incidence rate 12.1 p.m.p/year 2017-							-
2017- 2019 112 2054 9.3 Adults Gelgium 1152 glomerular final clinical diagnoses, FSGS incidence rate 12.1 p.m.p/year							_
Belgium Bilgium Belgium Bilgium Belgium Bilgium Belgium Bilgium Bilgium Bilgium Belgium Bilgium Bilgium Bilgium Bilgium Bilgium Belgium Bilgium B							
Belgium Bilgium Belgium Bilgium Belgium Bilgium Belgium Bilgium Bilgium Bilgium Belgium Bilgium Bilgium Bilgium Bilgium Bilgium Belgium Bilgium B							nephrology centers
Belgium							
Belgium Belgium Belgium Germany Belgium Belgium Belgium Belgium Belgium Germany Germany 1990-2013 1208 1208 Infirst period General				2054	9.3	Adults	(Belgium) 1152
Belgium Belgium Belgium Belgium Clinical diagnoses, FSGS incidence rate 12.1 p.m.p/year Diagnosis was often determined by results of genetic analysis Western France in the north of Brittany (Cotes d'Armor Depart ment) FSGS accounted 6.1% of all diagnoses (N=1,208). Over time FSGS was 4% General in first period in first period Clinical diagnoses, clinical diagnoses, clinical diagnoses, clinical diagnoses, and clinical diagnoses, clinical diagnoses, clinical diagnoses, are the p. SGS was 4% General clinical diagnoses, clinical diagnoses, clinical diagnoses, are the p. SGS was 4% General clinical diagnoses, clinical diagnoses, are the p. SGS was 4% General clinical diagnoses, clinical diagnoses, are the p. SGS was 4% General cl			2019 112				
FSGS incidence rate 12.1 p.m.p/year 2017- 2020 ¹¹³ 148 11.1% Children Diagnosis was often determined by results of genetic analysis Western France in the north of Brittany (Cotes d'Armor Depart ment) FSGS accounted 6.1% of all diagnoses (N=1,208). Over time FSGS was 4% in first period FSGS incidence rate 12.1 p.m.p/year Diagnosis was often determined by results of genetic analysis Western France in the north of Brittany (Cotes d'Armor Depart ment) A single center in Central Europe over a period of 24 years. 23 children (\leq 15 years) and 1,185		Belgium					
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		C					_
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$							rate 12.1
							p.m.p/year
France 1976 - 1990^{114} 480 10.6 General							Diagnosis was
France 1976 - 1990^{114} 480 10.6 General			2017-	140	11.107	Children	often determined
France 1976- 1990 114 480 10.6 General France in the north of Brittany (Cotes d'Armor Depart ment) FSGS accounted 6.1% of all diagnoses Germany 1990- 2013 18 1208 TSGS was 4% in first period General General Western France in the north of Brittany (Cotes d'Armor Depart ment) A single center in Central Europe over a period of 24 years. 23 children (≤15 years) and 1,185			2020 ¹¹³	146	11.1%	Children	by results of
France $\begin{array}{c} 1976-\\ 1990^{114} \end{array} \hspace{0.2cm} 480 \hspace{0.2cm} 10.6 \hspace{0.2cm} 10.6 \hspace{0.2cm} General \hspace{0.2cm} \begin{array}{c} \text{the north of} \\ \text{Brittany (Cotes} \\ \text{d'Armor Depart} \\ \text{ment)} \end{array}$ $\begin{array}{c} \text{FSGS accounted} \\ 6.1\% \text{ of all} \\ \text{diagnoses} \\ \text{over a period of} \\ \text{over a period of} \\ 24 \text{ years. } 23 \\ \text{children (\leq15} \\ \text{years) and } 1,185 \end{array}$							genetic analysis
France 1976 - 1990^{114} 480 10.6 General Brittany (Cotes d'Armor Depart ment) FSGS accounted 6.1% of all diagnoses 6.1% of all diagnoses 6.1% Over 6.1% 6.1% Over 6.1							Western France in
France 1990^{114} 480 10.6 General Brittany (Cotes d'Armor Depart ment) FSGS accounted 6.1% of all Central Europe diagnoses over a period of $(N=1,208)$. Over time FSGS was 4% General children (≤ 15 years) and 1,185			1076			General	the north of
Germany FSGS accounted 6.1% of all d'Armor Depart ment) A single center in Central Europe over a period of (N=1,208). Over time FSGS was 4% in first period General children (≤15 years) and 1,185		France		480	10.6		Brittany (Cotes
			1990***				d'Armor Depart
Germany $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$							ment)
Germany $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$					FSGS accounted		A single center in
Germany 1990- 2013 18 1208 (N=1,208). Over time FSGS was 4% in first period (N=1,208). Over time FSGS was 4% in first period 24 years. 23 children (≤15 years) and 1,185					6.1% of all		Central Europe
Germany 1990- 2013 18 1208 time FSGS was 4% General children (≤15 years) and 1,185					diagnoses		over a period of
Germany 2013 ¹⁸ 1208 time FSGS was 4% General children (≤15 years) and 1,185			1000		(N=1,208). Over		24 years. 23
in first period years) and 1,185		Germany		1208	time FSGS was 4%	General	children (≤15
(1990-1997), 10% adults. 706			2013 10		in first period		years) and 1,185
					(1990-1997), 10%		adults. 706
in second period (58.4%) was PGD.					in second period		(58.4%) was PGD.
(1998-2005) and Age averaged 50 ±					(1998-2005) and		Age averaged 50 ±

10% en 3rd period (2006-2013), being significant the trend in the	17.5 years, 63% male.
	į i
trend in the	
incidence of GSFS	
in the study period	
Primary FSGS was	
12%, secondary	
FSGS was 9%.	
FSGS had an	A single center
incidence of 11.2	study. The male-
pmp, with 43%	to-female ratio
2002- linked to an Adults	was 0.9, and the
2008 ¹¹⁵ underlying	rate of elderly
etiology.	persons aged 60
Secondary	years and more
glomerulonephritis	was 26.2%
had an incidence of	
17.5 pmp.	
	A tertiary referral
FSGS was 41% of	hospital. Average
Cyprus 2006- PGD, 12% primary Adults	age was 45.7
2015 116 and 29% secondary	years, 51% male;
form.	56% was PGD.
FSGS was 285	Nationwide renal
cases (13.2%) in	biopsy data in
all three-time	Lithuania.
Lithuania 1994- 2165 intervals. In 404 General	Average age was
2012 117 child patients	43.2 ± 20 years.
FSGS was 56 cases	Male: female ratio
(13.9%).	was 1.4:1.
	Average age was
2000-	$38.7 \pm 17.7 \text{ years},$
Estonia 2010 118 578 16.1 General	predominantly
	male; 45.4% PGD
1998-	One center. The
Portugal 228 7.9 Children	most common

					indication for
					kidney biopsy was
					nephrotic
					syndrome
					(42.9%). Primary
					glomerular
					diseases were
					found in 153 cases
				6 .	(67.1%) and FSGS
					corresponded to
					7.9% (18 cases)
					Six hospitals,
					predominantly
Finland	1976-	2057	2.0	Children	male. FSGS was
rillialid	2000^{120}	2037	3.9	Children	81 cases (3.9%)
					and, in patients
					<15 years was 3%.
					Norwegian Kidney
					Biopsy Registry
					(NKBR) and in the
					Norwegian Renal
Norway	1988-	575	8.17	Children	Registry (NRR).
Noiway	2021121	373	0.17	Ciliuren	Average age was
					10.7 (6.1 to 14.1)
	1				years, 313 (54.4%)
					were boys, FSGS
					was 47 patients.
					Mean age was 49
					± 17.8 years, 61%
					were male. PGD
United	1976-				was 907 patients
	1976- 2005 ¹²²	1844	5.7	Adults	and 52 was FSGS.
Kingdom	2003				Incidence de
					FSGS of 0.15
					php/year, per
					hundred thousand

						adult population
						per year.
						Average age was
						44 ± 20 year, 111
						were male, 33 of
		1974-	195	16.9	Adults	195 cases were
		2003 123	193	10.9	Adults	FSGS. Annual
						incidence for
						FSGS was 1.8 per
						100,000.
		1994-				281 PGD. FSGS
		2013 124	370	16	Adults	was 46 patients
		2013				with PGD.
						Average age was
					Adults	50.6 ± 16.7 years,
	United		2501	38.9		45.7% were
		2000- nited 2011 125 ates				women. FSGS was
						in 973 patients,
The						mean age in FSGS
Americas	States					group was 51.1
Americas						±16.2 years and
						56.9% were male.
		2001-				FSGS were in 435
		2005 126	1228	9.6	Adults	of 1228 adult
		2005				patients.
						A single center,
						patients >18 years.
						FSGS were 158
						patients. Mean age
		2004-	710	22.25	Adults	of the group with
		2014 127	, 10		1100105	FSGS was 54
						± 19.09 years, the
						ratio male: female
						was 1.72:1 in
						FSGS group.
	Brazil	1992-	582	28.8	General	9 AMICEN
	DIGE	2016 128	552	25.0	Conorui	(Minas Gerais

		1			Association of
					Nephrology
					Centers). Age
					means 35, 50.9%
					male sex. PGD
					was 75.3% of
					cases, FSGS was
					present in 126
					cases.
					Retrospective
					study, mean age of
					the general
					population was
	1993-	9617	24.6 in adults and	General	35.07 ± 18.65
	2007 129	9017	23.5% in Children	General	years, 51%
			, O'		female. (51.0%
					were PGD and
					FSGS was present
					in 1135 cases.
					Average age was
	1998- 2016 ¹³⁰	1151	43	Adults	$35.0 \pm 15.3, 41\%$
					male, 670 biopsies
					of native kidneys
					on patients.
					Paulista Registry
					of
					Glomerulopathies.
					1131 were PGD.
	1999-	2086	PGD in 29.7	Adults	Average age was
	2005 131				34.5 ± 14.6. PGD
					were more
					frequent in males
					(55.1%).
	2000- 2018 ¹³²	1051			Average age was
			PGD in 37.3% and 10% secondary FSGS.		44.9 ± 16.1 years.
				Adults	Female 52.9%,
	2010				FSGS in 60.3%.
					1 505 III 00.5 /0.

					Temporal
					variation across
					the three time
	2000-				periods showed a
	2018 133	1057	37.3%	Adults	statistically
	2018				-
					significant
					reduction in FSGS
					over time
				<u> </u>	FSGS was present
					in 288 biopsies
	1998-	1040	34.8 in adults and	Adults	(109 were male);
	2007 23		28.7% in children.		in children were
					76 patients (male
					were 48 patients).
					Median age of
		1412	Primary FSGS in 20.6	General	patients was 26
					years; 56.7% were
					males, 291 had the
	1998-				confirmed
	2009 134				diagnosis of
					primary FSGS, 74
Color	mbia				patients (25.4%)
	(were < 15 years of
					age.
					Over 18 years old.
					Male: female ratio
	2003-	9911	20.1	Adults	was 42.6%:
	2015 135	<i>)</i>	20.1	rauits	57.4%. 1992
					patients had
					FSGS.
					Average age was
	2007			Children	11±4.3 years, 58%
	2007-	241	11.6		female. FSGS was
	2017 136				present in 28
					patients.
				1	

		2000				The mean age was
		2008- 2018 ¹³⁷	871	19	Adults	39 ± 14 years,
		2018137				67% female.
						Single second
						level hospital
						center.
						Retrospective
						analysis, Average
		2003-				age was 32.6 ±
	Mexico 138	2011	163	47	Adults	13.3, 55% were
		2011				female. In FSGS
						group male:
						female ratio was
						1.4:1 and average
						age was 25.9 ±
						10.4 years.
		da 1985- 2014 ¹³⁹	6434	The relative	Adults	
				frequencies of		Mean age 47.9 ±
	Canada			FSGS was 13.73%		
				(1985-1989),		
				16.13% (1990-		19.8 years, 58%
	Curacu			1994), 17.93%		were male.
				(1995-1999),		
				15.99% (2000-		
				2004) and 17.9%		
				(210-2014).		
	Chile	1999- 2020 ¹⁴⁰	550	14.1	Adults	63.5% females, Mean age at diagnosis was 47.8 ± 18.2 years.
				FSGS was the		Incidence FSGS
				most frequent PG		was 6.4 per
				in 29.3% and		million population
	Итионом	1980- 2003 ¹⁴¹	2058	decreased from	Adults	(pmp). Average
	Oruguay		2030	36.3% in 1995-	Muits	age was 39.1 ±
				1999 period to		19.6 years, males
				19.1% in 2000-		was >50% on all
				2003 period.		periods.

	I					ECCC
						FSGS was present
						in 144 patients 171
						were idiopathic. In
				Global FSGS was		addition, 101 cases
		1985-		13.9 and 15.8 in		were analyzed in
	Peru	1995 ¹⁴²	1263	prospective	Adults	prospective and
				patients.		FSGS was present
				1		in 16 patients with
						mean age 31
						years, sex
						male/female 5/11.
						Between 1997 and
						2001 FSGS
						predominated in
				31.8 between		56 patients, and
		1997-	162	1997-2001 and 43	Children	between 2006 and
	Nigeria	2013 143		between 2006-		2013, native
				2013		kidney biopsy of
						106 children,
						FSGS was in 46
				>		patients.
			165	27.8	General	Single center.
		2002-				Average age was
Africa		2011 144				15.4 ± 12 years,
Airica						64.8% male.
	Egypt	2003-	924	21.21 of PGD	General	Mean age was
	Lgypt	2008 145	724	21.21 011 0D	General	26.5 ± 14.6 years.
						A single-center
						renal biopsy
	South	2000-	1284	10.5	Adults	database. Average
	Africa	2009 146	1204	10.3	Adults	age was 36.8 ± 14
						years, 54% were
						female.
	M					A single center
		2000- 2007 ¹⁴⁷	161	5.0	Adults	renal biopsy
	Morocco			5.9		database. PGD
						were in 84
	<u> </u>	İ	<u> </u>	L	<u>I</u>	

		1	<u> </u>			notionto Avonoco
						patients. Average
						age was 40.4 ± 15
						years, 101 males,
						FSGS was in 8
						patients.
						Mean age was
	Sudan	2002-	321	13.7	Children	8.71, 60.2% were
	Sudan	2007 148	321	13.7	Cilidren	male. FSGS was
						in 44 patients.
						Age range non-
						indigenous
			653 renal	In 249 non-		42.5±16.6 years,
			biopsies	indigenous patients		aboriginal
			on	FSGS was 16.1%,		remote/very
			indigenous	in aboriginal non-		remote (n=455)
		1982- 2005 ¹⁴⁹	people and	remote 12%, in	Adults	age was
			249	Torres Strait		39.2±13.9. Non-
			biopsies	Islander was 9.5%,		indigenous 40.2%
			non-	in aboriginal		female, and
			indigenous	remote/very remote		aboriginal
			patients.	FSGS was 19.6%.		remote/very
						remote 55.4%
	Australia					female.
Oceania				FSGS was relative		
				frequency of		
				20.9% in patients		
				with nephrotic		
				syndrome and		Average age was
		2002-		6.4% in nephritic		48 ± 17 years.
		2011 17	3697	syndrome,	Adults	Male was $\sim 60\%$,
				undefined renal		FSGS was in 1.02
				dysfunction 15.9%		php/yr.
				and nephrotic		
				proteinuria was		
				35.3% of cases.		
	New	1986-				FSGS was present
	Caledonia	1993 ¹⁵⁰	181	20.4	Adults	in 37 cases.
		l				

FSGS: Focal Segmental Glomerulosclerosis; JNSCS Japan Nephrotic Syndrome Cohort Study; J-RBR: Japan Renal Biopsy Registry; PGD: Primary glomerular diseases; NS: Nephrotic Syndrome SRNS: Steroid-resistant nephrotic syndrome