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Review

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



Una aproximación a la prevalencia de la glomeruloesclerosis focal y segmentaria: una revisión sistemática de la literatura mundial durante los últimos 32 años

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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.

RESUMEN

Palabras clave: Glomeruloesclerosis focal y segmentaria Glomerulonefritis Biopsia renal Prevalencia Antecedentes: La glomeruloesclerosis focal y segmentaria (GEFS) es una lesión histopatológica caracterizada por la cicatrización en secciones específicas de algunos glomérulos, acompañada de lesión de podocitos. A nivel mundial, la prevalencia de la GEFS y sus tendencias temporales no han sido suficientemente estudiadas. Sin embargo, algunos informes sugieren un aumento en la frecuencia de la GEFS en las últimas décadas. Comprender la epidemiología de la GEFS es crucial para que los médicos mejoren el diagnóstico y el tratamiento.

Objetivo: Este estudio evalúa críticamente las tendencias de prevalencia mundial de FSGS en los últimos 32 años (1992-2024), destacando las variaciones entre países a través de una revisión sistemática.

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Métodos: Se realizó una búsqueda sistemática en Medline, Embase y ScienceDirect para identificar estudios relevantes. La fiabilidad de los datos de prevalencia se evaluó mediante una evaluación crítica de publicaciones seleccionadas.

Resultados: La prevalencia de la GEFS varía significativamente de una región a otra. Los países de Asia Oriental tienen una prevalencia relativamente baja, con una media de alrededor del 7%. En contraste, los países del sur de Asia, Medio Oriente y las Américas tienen una mayor prevalencia, con un 18%. Los países europeos muestran una prevalencia intermedia de alrededor del 11%. Los países africanos no muestran un patrón claro, con tasas de prevalencia alta y bajas en diferentes países.

Conclusiones: La prevalencia de la GEFS difiere según la región geográfica y la etnia. Mientras que los países del sur de Asia han mantenido una prevalencia sistemáticamente baja, otras regiones han experimentado un aumento de los casos de GEFS a lo largo del tiempo. Este estudio mejora la comprensión de los patrones globales de FSGS, proporcionando información epidemiológica valiosa para médicos e investigadores.

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. 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.

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.

Methods

The search strategy followed the preferred reporting items for systematic reviews and meta-analysis (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.

Results

The initial literature search identified 1047 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

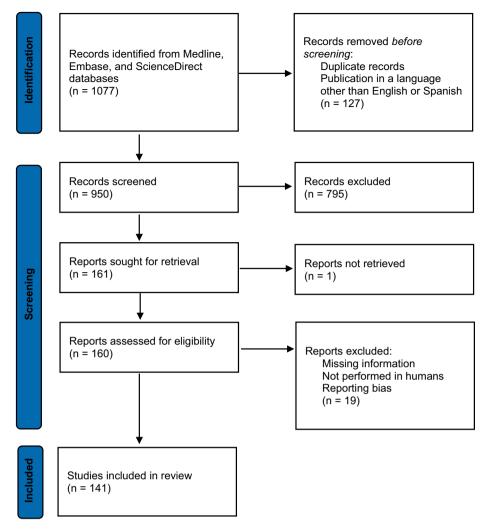


Fig. 1. PRISMA flowchart depicting the study selection process.

excluded at each stage, is presented in Fig. 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.

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.

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

Continent	Country	Period and reference	No. biopsies	Prevalence (%)	Population	Description
Asia	China	1979–2002 ²⁹	13,519	6	Adults. A single unit in China and 57.3% were male.	Average age was 32.7 \pm 12.2 (9–83) years. Biopsies with PGD 9278, FSGS was present in 557 cases.
		1987–2012 ³⁰	11,618	5.82	General; All patients were Asian, 6646 male cases and 4972 female cases (1.33:1).	PGD 8209 cases, median age of 35 years (range 3–85 years old). Among the PGD FSGS was present in 458 patients of 8209.
		1993–2017 ³¹	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 \geq 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 (2000–2004) to 5.57% in period 3 (2005–2009), but then it decreased to 3.09% in period 4 (201–2014).	Adults	Ten hospitals in China. Average age was 35.2 years, 2254 patients (45.7%) were men. PGD were 81.55%, FSGS was 146 patients (3.63%), 85 male and 61 female.
		2001–2015 ³³	10,779	2.5	General	A single center, patients aged (\geq 15 years old. Average age was 40 \pm 14.8 years old
		2001–2019 ³⁴	9448	FSGS was in 477 patients (5.05%), group 1 (01/01/2001 to 12/21/2004) was 51 patients (5.06%), group 2 (01/01/2005 to 12/31/2009) was 109 patients (6.04%), group 3 (01/01/2010 to 12/31/2014) was 205 patients (6.26%), and group 4 (01/01/2015 to 12/31/2019) was 112 patients (3.33%).	Adults	Average age was 41.2 years for males and 41.60 years or females. 60.39% male.
		2003–2014 ³⁵	40,759	FSGS was 6% in period 1979–2002 and changes frequencies at 7.34% to period of 2003–2014.	Adults	Patients $>$ 14 years. Renal Biopsy Registry of the National Clinical Research Center of Kidney Diseases. Average age was 36.59 ± 14.12 years. 52% male. PGD was 67.1% of cases.
		2004–2014 ³⁶	7962	FSGS was 6% in male and 4% in female patients. 5% in patients to 0–12 year and 6% in 13–18 year. FSGS showed a significant decreasing trend, starting at 14% in the first period (2004–2007) and decreasing to 6% and 4% in the second (2008–2011) and third periods (2012–2014).	Children	Average age was 13.5 ± 4.1 years, 64% boys. Included 115 hospitals.
		2007–2016 ³⁷	2725	2.68	Adults	One hospital in northeast China. Only >14 years. Average age was 41.2 \pm 15.1 years. FSGS was 73 patients.
		2008–2018 ³⁸	10,996	8	General	A single center, retrospective study of native kidney biopsies, PGD was 69.42%, with 8% for GSFS.
		2008–2017 ³⁹	4910	3.1	General	3593 cases of PGD (73.2%). Average age was 42.6 \pm 15.7 years (range: 7 to 84 years), 2629 males and 2281 females (ratio: 1.15:1).
		2008–2013 ⁴⁰	3722	8.78	Adults	Shandong Province of China. FSGS patients average was 37.9 ± 12.5 years FSGS frequency was 327 patients.
		2009–2018 ⁴¹	35,783	FSGS frequency was in 850 patients (2.45%). FSGS had distribution peaks in the 20–39-year age group. In elderly patients FSGS was 2.92% and in children (-14 years) FSGS was 3.32%.	General	Adults 54.27% male patients, ratio male. Female 1.19:1, 31,256 were adults, with age of adults 40.8 \pm 15.2 years.

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

Table 1

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

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Continent	Country	Period and reference	No. biopsies	Prevalence (%)	Population	Description
		2011–2020 ⁴²	9310	14.97	Adults	One center, two periods 2011–2015 and 2016–2020. PGD were in 66.93% cases (6166 cases), among these FSGS was 14.97% patient In patients with age 14–24 years FSGS was 14.04%, in 25–44 year 13.39% and 45–59 years 18.33%.
		2014–2016 ⁴³	1445	21.6	Adults	Included 17 centers, of PGD FSGS was 21.6% cases (133). Average at was 50.4 ± 17.7 years, 55.6% male, 14.3% had diabetes and 50.4 had hypertension.
		2014–2018 ⁴⁴	7917	4.3	Adults	Nationwide cross-sectional survey of kidney biopsies in China. 59.6 male. FSGS was in 331 patients.
		2015–2017 ⁴⁵	62,569	FSGS was in 2526 (4.4%) of patients, 199 (4.7%) pediatrics and 2327 (4.4%) in adult patients	General	Retrospective study, including 56,880 native biopsies at 1211 hospitals across China. Children were 4274 patients y adults 52,606 patients.
	India	1990–2008 ⁴⁶	1849	15.3	General	Average age was 32.2 ± 18.3 years. 1091 males and 758 females. FSFGS was 195 of 1278 patients, median age 25 years. Male 2.25 1 Female
		1997–2013 ⁴⁷	423	25.8	Children	A single center. Average age was 10.4 ± 4.5 years, 57.9% male. PG was 360 patients, and of these FSGS was 109 patients.
		2006–2016 ⁴⁸	3257	FSGS was the most common PGD accounting for 18.2%.	Adults	A single center study in northern India. Average age was 33.2 ± 14 . years, 61.9% male.
		2007-2018 ⁴⁹	254	12.2	Children	Median age was 15 years, 57% female.
		2008-2013 ⁵⁰	335	8.02	Children	Average age was 7.91 ± 3 years, 68.1% males. PGD was the most common in 81.8%, and FSGS was 8.02%.
		2009–2016 ⁵¹	270	13.7	Adults	Retrospective study. Average age was 36.92 years. FSGS was 37 out 270) most common PGD.
		$2010 – 2012^{52}$	666	22.58	Adults	Average age was 28 \pm 14.6 years. Of 527 cases, primary FSGS wa 119 patients.
		2016-2021 ⁵³	409	17.4	Children	353 PGD, FSGS was present in 71 patients.
		2017–2020 ⁵⁴	347	17.02	Adults	Mean age was 41.4 ± 15.7 years, 58.5% were males. NS 36.3% of cases; 69% was PGD, FSGS was 32 patients.
	Japan	2007–2008 ⁵⁵	2126	3.62	Adults	Japan Renal Biopsy Registry. Prospective registry system, J-RBR, 23 renal centers. Average age was 44.4 ± 21 years, 1281 male. FSC was present in 77 patients.
		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 45 patients.
		2007–2011 ⁵⁷	2802	FSGS present in 99 patients from 1259 patients in elderly (≥65 years old) and 83 patients (1.7%) in control group (20–64 years) of 5021 cases.	Adults	J-RBR of the Japanese Society of Nephrology included 1596 male Renal disease in the elderly (age ≥65 years old) and very elderly (ag ≥80 years old) Japanese.
		2007–2016 ⁵⁸	1409	3	Adults	JRBR. Average age was 47 years, 59.4% were male. The annual incidence of FSGS accounted for 3.5–4.5% Percentage of primary FSGS were constant at approximately 3% of all during 2007 to 201
		2007–2017 ⁵⁹	32,254	3.4 in adults and 3.7 in pediatrics	General	Japan Renal Biopsy Registry (J-RBR) 2007–2017. 973 patient adul with FSGS and 129 pediatrics patients with FSGS.
		2009–2010 ⁶⁰	8697	5.3	Adults	JRBR. Predominantly males 53.6% in 2009 and 54.9% in 2010, included 4016 biopsies in 2009 and 4681 cases in 2010, average at (years) were 46.7 ± 19.9 and 46.7 ± 20.6 respectively. From 7034 FSGS was present in 378 patients.
		2015-2018 ²²	6036	10.85	Adults	JNSCS patients with primary NS; FSGS was 655 patients, age 39 year 57.5% males.
	Pakistan	1997–2013 ⁶¹	423	25.8	Children	Average age was 10.48 \pm 4.58 years, 57.9% were males (245). FSC in 109 patients.
		1998–2005 ²⁵	415	Primary FSGS in 18.3% of patients.	Children	Pediatric kidney biopsies between 3–15 years, male: female ratio w 1.6:1.

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

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Continent	Country	Period and reference	No. biopsies	Prevalence (%)	Population	Description
		2010-2015 ²⁶	108	25.9	Children	A tertiary care hospital. Average age in FSGS patients was 7.0 ± 4.2 years, 56.5% were male, FSGS was present in 28 patients.
		2011-2020 ⁶²	307	Primary FSGS in 40.4% (124 patients)	Children	. FSGS cases, mean age was 8.8 ± 3.0 years while most of the children, 70 (56.5%) were above 10 years of age; 64 cases were male.
	South Korea	1973–1995 ⁶³	2097	FSGS was present in 154, Adults 97 (4.6%) and 57 Children (4%).	General	82.5% PGD. Male: female ratio was 1.5:1.
		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, male-to-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 \pm 16.5 years, 56% were male.
	Iran	2006-2018 ⁶⁷	2975	15.9	Adults	The mean age of the patients was 27.4 years old; 51.6% were male.
		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 glomerulonephritis (24.2%).	Adults	A Single Center. Average age was 33.9 ± 17.5 years, 58% men.
		$2007 - 2018^{70}$	2975	15.9%	Adults	A Single Center. 51.6% was male. The mean age was 27.4 years.
	Saudi Arabia	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.
		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.
		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.
		1991–2010 ⁷⁵	3892	16.46	Children	Turkish Society of Nephrology Registry System (TSNRS), 2438 were glomerular diseases The age range was 0 to 15 years. Male/female ratio was 1:1, FSGS was 641 patients.
		2009–2019 ⁷⁶	3875	21.9	Adults	A multicenter study by the Turkish Society of Nephrology Glomerular Diseases Working Group, 47 centers of PGD of Turkish Society of Nephrology Glomerular Diseases (TSN-GOLD) Working Group. Average age was 41.5 ± 14.9 years, 56.3% were male.
	Jordan	2010–2016 ⁷⁷	99	23	Children	Retrospective study of FSGS, a tertiary care hospital. Average age 3.71 ± 2.59 years, 66% were male.
		2013-2020 ⁷⁸	106	11.32	Adults	A Single-Center. Average age was 34 \pm 12.7 years, females were 53.7%
	Nepal	2014–2016 ⁷⁹	175	18.28% (18 patients) had FSGS.	Adults	A single center. Tertiary care center. Average age was 35.3 ± 13.5 years. 59.4% female, ratio female: male ratio of 1.5. The mean age of patients with FSGS was 36.89 years.
		2001-2007 ⁸⁰	137	8	General	Average age was 30.6 years for males and 32.9 years for females. The male to female ratio was 1.6: 1
		$2022 - 2022^{81}$	213	Primary FSGS in 10.33	Adults	One Center
	Lebanon	2003–2007 ⁸²	1048	In Age \leq 15 FSGS was 6.6%, in 15 $<$ Age \leq 60 FSGA was 13.9% and age $>$ 60 FSGS was 12.1%.	General	Average age was 36.76 ± 20 years (range 1–84), 54.4% were male, 20% pediatric population, 67.3% young adult group and 12.7% elderly group.

Table 1 (Continued)
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
		2014–2015 ⁸³	144	Children were FSGS (5.7%) 31 adult patients were FSGS (28.4%)	General	35 children (mean age 11 ± 5.6 years) and 109 adults (mean age 41.6 ± 16.5 years). 74.2% were male in children group and 45.8% were male in adult patients,
	United Arab Emirates	1978–1996 ⁸⁴	490	18.3	Adults	Data from the United Arab Emirates Renal Diseases Registry. 378 PGD. Adults aged 14–66 years. FSGS were 69 patients.
	Taiwan	2014–2016 ⁸⁵	1445	FSGS were in 133 of 599 PGD patients (25%),	Adults	National Renal Biopsy Registry with 17 medical centers. Average age was 48.4 ± 16.6 years, 53.8% cases male samples. Average age 50.4 ± 17.1 years, 55.6% male in FSGS group.
	Thailand	1983-2005 ⁸⁶	3355	2.8	Adults	A single center, FSGS was 61 patients.
	Oman	2011–2015 ⁸⁷	596	Primary FSGS in 18.3% patients.	Children	Pediatric kidney biopsies between (3–15 years), male: female ratio was 1.6:1.
	Iraq	2012-2013 ⁸⁸	662	22	Adults	Average age was $27.3 \pm 17.6, 53\%$ were male. FSGS was 135 patients.
	Bangladesh	2008–2009 ⁸⁹	95	11.58	Adults	Average age was 30.29 years, 60% female; male to female ratio of 1:1.5.
	Sri Lanka	2018–2019 ⁹⁰	140	22.1	Adults	One center, 55.7% females. Mean age was 46 ± 15.3 years. FSGS in 31 patients.
	Kuwait	2013–2018 ⁹¹	356	21.9	General	Report of four public hospitals, 356 adult native kidney biopsies, average age was 39.8 (12 to 90 years) years, 61.2% were male. FSGS came third place with 78 (21.9%) of the cases, 31 patients were less than 18 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	3282 biopsies with PGD	11.88	General	Average age was 47.9 \pm 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 \pm 12.2 years to 47.0 \pm 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 (range 0.6–24 years), 43.2% female.
		1987–1993 ²⁰	13,835	FSGS was 11.8% of glomerulonephritis. The annual frequency of PGD to FSGS was 10.4% in 1993, 14.4% in 1990.	Adults	The Italian Group of Renal Immunopathology. Male sex was predominant in PGD (65%).
		1996–2000 ⁹⁵	14,607	FSGS was 16.9% of PGD with NS	Adults	Italian Immunopathology Group, Date from 128 rental units in Italy were reported. PGD were in 6990 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
		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.
		2011–2019 ⁹⁹	1101	4.8% of PGD (2.9% of 442 cases)	Adults	Biopsy reports were divided into 3 periods, 320 from 1994 to 2004 (period 1), 239 from 2005 to 2010 (period 2) and 442 biopsies between 2011 and 2019 (period 3). Mean age of the renal biopsy population during period 3 was 39.2 ± 13.8 years, 65.2% were male. PGD was 59.5% of the cases.
	Poland	1990–2010 ¹⁰⁰	746	FSGS was 58 of 607 PGD patients (9.5%).	Adults	Adults (> 18 years), in a single tertiary nephrology center serving an area of Central Poland. 607 PGD. Average age was $40.5 \pm 8\ 20.8$ years, 411 male patients.

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

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Continent	Country	Period and reference	No. biopsies	Prevalence (%)	Population	Description
		2009–2014 ¹⁰¹	8843	15	General	Polish Society of Nephrology. A total of 1939 (21%) biopsies were performed in patients <18 years of age, 6394 (68.7%) in those 18–64 and 955 (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.
	Spain	1994–1999 ¹⁰²	7016	In children FSGS was 15.2% and 10.8% in adult patients; in the elderly FSGS was 6%.	General	93 medical renal units. Median (range) < 15 years (487), 15–65 years (4827) and > 65 years (1510). Male/female ratio in children of 1.2, in adults of 1.5 and in the elderly of 1.4.
		1994-2019 ¹⁰³	18,852	9	Adults	The age range was 15 to 65 years. Male/female ratio was 1.5.
	Sweden	2014-2029 ¹⁰⁴	913	8.4	Adults	FSGS was present in 77 patients.
	Czech Republic	1994–2000 ¹⁰⁵	3294 biopsies in adults and 710 biopsies in <15 years	10.8	General	Czech Registry of Renal Biopsies included 28 centers. Mean age 10 years for children and 42 years for adults, 57.9% males.
		1994–2011 ¹⁰⁶	10,472	12.6	General	31 centers, 57.8% male. Mean age for children was 10 years and for adults 44.5 years FSGS incidence was 3.9 p.m.p/year, mean age for FSGS group was 40 years, 56.8% males.
	Denmark	1985–1997 ¹⁰⁷	2380	13.66	Adults	Danish Renal Biopsy Register (DANYBIR). Average age was 42.6 ± 20.2 years. FSGS was 325 patients. Incidence 5.7 pmp/year, age FSGS group was 43.4 ± 19 and 34% were female.
		1985–2014 ¹⁰⁸	5043	7.93	Adults	Danish Renal Biopsy Registry and Patobank registries. FSGS was 400 patients of 5043 biopsies.
	Serbia	1987–2006 ¹⁰⁹	1626	FSGS was 15.1% causes of nephrotic syndrome among 872 native kidney diseases.	Adults	Average age was 39.1 \pm 13.8 years, 51.2% were male.
		2001–2010 ¹¹⁰	150	20.9	General	A single center. Mean age was 11.5 years, 56% were female patients. PGD was 57.4%.
	Croatia	1996–2012 ¹¹¹	922	15.8	Adults	Average age was 48 years (range 16–84 years), patients were \geq 16 years old.
	Belgium	2017–2019 ¹¹²	2054	9.3	Adults	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
		$2017 - 2020^{113}$	148	11.1%	Children	Diagnosis was often determined by results of genetic analysis
	France	1976-1990 ¹¹⁴	480	10.6	General	Western France in the north of Brittany (Cotes d'Armor Department)
	Germany	1990–2013 ¹⁸	1208	FSGS accounted 6.1% of all diagnoses (<i>N</i> = 1208). Over time FSGS was 4% in first period (1990–1997), 10% in second period (1998–2005) and 10% en 3rd period (2006–2013), being significant the trend in the incidence of GSFS in the study period	General	A single center in Central Europe over a period of 24 years. 23 children (\leq 15 years) and 1185 adults. 706 (58.4%) was PGD. Age averaged 50 \pm 17.5 years, 63% male.
		2002–2008 ¹¹⁵	222	Primary FSGS was 12%, secondary FSGS was 9%. FSGS had an incidence of 11.2 pmp, with 43% linked to an underlying etiology. Secondary glomerulonephritis had an incidence of 17.5 pmp.	Adults	A single center study. The male-to-female ratio was 0.9, and the rate of elderly persons aged 60 years and more was 26.2%
	Cyprus	2006–2015 ¹¹⁶	153	FSGS was 41% of PGD, 12% primary and 29% secondary form.	Adults	A tertiary referral hospital. Average age was 45.7 years, 51% male; 56% was PGD.
	Lithuania	1994–2012 ¹¹⁷	2165	FSGS was 285 cases (13.2%) in all three-time intervals. In 404 child patients FSGS was 56 cases (13.9%).	General	Nationwide renal biopsy data in Lithuania. Average age was 43.2 ± 20 years. Male: female ratio was 1.4:1.
	Estonia	$2000 – 2010^{118}$	578	16.1	General	Average age was 38.7 \pm 17.7 years, predominantly male; 45.4% PGD

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

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Continent	Country	Period and reference	No. biopsies	Prevalence (%)	Population	Description
	Portugal	1998-2021 ¹¹⁹	228	7.9	Children	One center. The most common indication for kidney biopsy was
						nephrotic syndrome (42.9%). Primary glomerular diseases were
	r:-11	1076 0000120	0057	2.0	Ob this are	found in 153 cases (67.1%) and FSGS corresponded to 7.9% (18 cases
	Finland	1976–2000 ¹²⁰	2057	3.9	Children	Six hospitals, predominantly male. FSGS was 81 cases (3.9%) and, in patients <15 years was 3%.
	Norway	1988-2021 ¹²¹	575	8.17	Children	Norwegian Kidney Biopsy Registry (NKBR) and in the Norwegian
						Renal Registry (NRR). Average age was 10.7 (6.1 to 14.1) years, 31 (54.4%) were boys, FSGS was 47 patients.
	United Kingdom	1976-2005 ¹²²	1844	5.7	Adults	Mean age was 49 ± 17.8 years, 61% were male. PGD was 907 patient
						and 52 was FSGS. Incidence de FSGS of 0.15 php/year, per hundre
						thousand adult population per year.
The Americas	United States	1974-2003 ¹²³	195	16.9	Adults	Average age was 44 \pm 20 year, 111 were male, 33 of 195 cases were
						FSGS. Annual incidence for FSGS was 1.8 per 100,000.
		1994–2013 ¹²⁴	370	16	Adults	281 PGD. FSGS was 46 patients with PGD.
		2000-2011 ¹²⁵	2501	38.9	Adults	Average age was 50.6 ± 16.7 years, 45.7% were women. FSGS was i
						973 patients, mean age in FSGS group was 51.1 ± 16.2 years and 56.9% were male.
		2001-2005 ¹²⁶	1228	9.6	Adults	FSGS were in 435 of 1228 adult patients.
		2001–2003 2004–2014 ¹²⁷	710	22.25	Adults	A single center, patients > 18 years. FSGS were 158 patients. Mea
			,			age of the group with FSGS was 54 ± 19.09 years, the ratio male:
						female was 1.72:1 in FSGS group.
	Brazil	1992-2016 ¹²⁸	582	28.8	General	9 AMICEN (Minas Gerais Association of Nephrology Centers). Age
						means 35, 50.9% male sex. PGD was 75.3% of cases, FSGS was present in 126 cases.
		1993-2007 ¹²⁹	9617	24.6 in adults and 23.5% in Children	General	Retrospective study, mean age of the general population was
						35.07 ± 18.65 years, 51% female. (51.0% were PGD and FSGS wa present in 1135 cases.
		1998-2016 ¹³⁰	1151	43	Adults	Average age was 35.0 ± 15.3 , 41% male, 670 biopsies of native
						kidneys on patients.
		1999–2005 ¹³¹	2086	PGD in 29.7	Adults	Paulista Registry of Glomerulopathies. 1131 were PGD. Average ag
		0000 0010132	1051	DCD: 07.00/ 1.100/ 1	4.1.1.	was 34.5 ± 14.6 . PGD were more frequent in males (55.1%).
		2000–2018 ¹³²	1051	PGD in 37.3% and 10% secondary FSGS.	Adults	Average age was 44.9 ± 16.1 years. Female 52.9%, FSGS in 60.3%
		2000-2018 ¹³³	1057	37.3%	Adults	Temporal variation across the three time periods showed a
	Colombia	1998-2007 ²³	1040	34.8 in adults and 28.7% in children.	Adults	statistically significant reduction in FSGS over time FSGS was present in 288 biopsies (109 were male); in children were
	Colollibia	1996–2007	1040	34.8 III addits and 26.7% III children.	Addits	76 patients (male were 48 patients).
		1998-2009 ¹³⁴	1412	Primary FSGS in 20.6	General	Median age of patients was 26 years; 56.7% were males, 291 had th
				, , , , , , , , , , , , , , , , , , , ,		confirmed diagnosis of primary FSGS, 74 patients (25.4%) were <1
						years of age.
		2003–2015 ¹³⁵	9911	20.1	Adults	Over 18 years old. Male: female ratio was 42.6%: 57.4%.
		196				1992 patients had FSGS.
		2007–2017 ¹³⁶	241	11.6	Children	Average age was 11 ± 4.3 years, 58% female. FSGS was present in
		2008-2018 ¹³⁷	871	19	Adults	28 patients. The mean age was 39 ± 14 years, 67% female.
	Mexico	2003–2011 ¹³⁸	163	47	Adults	Single second level hospital center. Retrospective analysis, Average
						age was 32.6 ± 13.3 , 55% were female. In FSGS group male: female.
						ratio was 1.4:1 and average age was 25.9 \pm 10.4 years.
	Canada	1985-2014 ¹³⁹	6434	The relative frequencies of FSGS was	Adults	Mean age 47.9 ± 19.8 years, 58% were male.
				13.73% (1985–1989), 16.13% (1990–		
				1994), 17.93% (1995–1999), 15.99%		
	Chile	1999-2020 ¹⁴⁰	550	(2000–2004) and 17.9% (210–2014). 14.1	Adults	63.5% females, Mean age at diagnosis was 47.8 \pm 18.2 years.
	CHILE	1777-2020	330	17.1	riduits	03.370 Temates, inean age at diagnosis was 47.0 ± 16.2 years.

Nefrologia 45 (2025)

Table 1 (Continued)
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
	Uruguay	1980–2003 ¹⁴¹	2058	FSGS was the most frequent PG in 29.3% and decreased from 36.3% in 1995–1999 period to 19.1% in 2000–2003 period.	Adults	Incidence FSGS was 6.4 per million population (pmp). Average age was 39.1 \pm 19.6 years, males was >50% on all periods.
	Peru	1985–1995 ¹⁴²	1263	Global FSGS was 13.9 and 15.8 in prospective patients.	Adults	FSGS was present in 144 patients 171 were idiopathic. In addition, 101 cases were analyzed in prospective and FSGS was present in 16 patients with mean age 31 years, sex male/female 5/11.
Africa	Nigeria	1997–2013 ¹⁴³	162	31.8 between 1997–2001 and 43 between 2006–2013	Children	Between 1997 and 2001 FSGS predominated in 56 patients, and between 2006 and 2013, native kidney biopsy of 106 children, FSGS was in 46 patients.
		2002-2011 ¹⁴⁴	165	27.8	General	Single center. Average age was 15.4 ± 12 years, 64.8% male.
	Egypt	2003-2008 ¹⁴⁵	924	21.21 of PGD	General	Mean age was 26.5 ± 14.6 years.
	South Africa	2000–2009 ¹⁴⁶	1284	10.5	Adults	A single-center renal biopsy database. Average age was 36.8 ± 14 years, 54% were female.
	Morocco	2000-2007 ¹⁴⁷	161	5.9	Adults	A single center renal biopsy database. PGD were in 84 patients. Average age was 40.4 ± 15 years, 101 males, FSGS was in 8 patients.
	Sudan	$2002 - 2007^{148}$	321	13.7	Children	Mean age was 8.71, 60.2% were male. FSGS was in 44 patients.
Oceania	Australia	1982–2005 ¹⁴⁹	653 renal biopsies on indigenous people and 249 biopsies non- indigenous patients.	In 249 non-indigenous patients FSGS was 16.1%, in aboriginal non-remote 12%, in Torres Strait Islander was 9.5%, in aboriginal remote/very remote FSGS was 19.6%.	Adults	Age range non-indigenous 42.5 ± 16.6 years, aboriginal remote/very remote ($n=455$) age was 39.2 ± 13.9 . Non-indigenous 40.2% female, and aboriginal remote/very remote 55.4% female.
		2002–2011 ¹⁷	3697	FSGS was relative frequency of 20.9% in patients with nephrotic syndrome and 6.4% in nephritic syndrome, undefined renal dysfunction 15.9% and nephrotic proteinuria was 35.3% of cases.	Adults	Average age was 48 \pm 17 years. Male was ~60%, FSGS was in 1.02 php/yr.
	New Caledonia	1986-1993 ¹⁵⁰	181	20.4	Adults	FSGS was present in 37 cases.

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.

(>14 years old), for whom more reports are available, show a slightly higher mean prevalence of approximately 7.06%.

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 8325 biopsies analyzed, with values ranging from 8.02% to 22.58%. The number of samples analyzed per study ranged from 65 to 3257. 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%.

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 2802. 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%.

Pakistan

The six studies conducted in Pakistan analyzed a total of 1375 kidney biopsies over a period from 1997 to 2021. Most reports focus on the pediatric population (≤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%.

South Korea

The South Korean population studies spanned from 1973 to 2013 and analyzed a total of 6657 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%.

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 4803 biopsy samples.

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.

Turkey

In Turkey, the mean prevalence of FSGS was approximately 15.22%, based on an analysis of 8381 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%.

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%.

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.

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%.

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.

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 2127 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.

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.

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.

United States

In the United States, the prevalence of FSGS was relatively high, with a mean of 20.13% according to six studies analyzing 5173 biopsies collected between 1974 and 2020, with some overlap

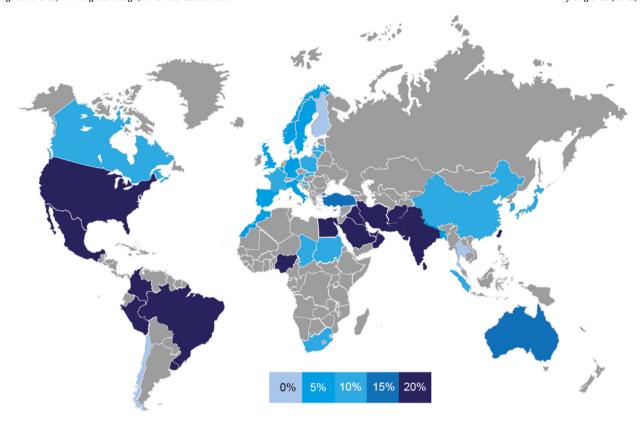


Fig. 2. Prevalence of FSGS in the studied countries. A blue color scale represents prevalence, with darker shades indicating higher rates. Countries shown in gray lack available prevalence data.

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 1994–2003 and 2004–2013.

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.

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.

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.

Oceania

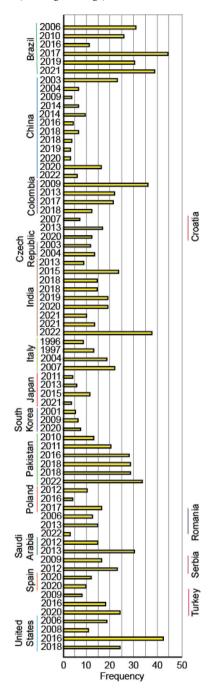
In Oceania, the prevalence of FSGS is relatively high, averaging approximately 18.53%. This estimate is based on an analysis of 4531 samples collected in Australia and New Caledonia between 1982 and 2011.

Discussion

The prevalence of FSGS varies significantly from region to region (Fig. 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%). 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 (Fig. 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



 $\textbf{Fig. 3.} \ \, \textbf{Trends in prevalence of FSGS over time in selected countries.}$

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 high-resource 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 and 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 to 11.6% in Italy, 18.3 to 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.

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.

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Conflicts of interest

The authors declare no conflict of interest.

Data availability statement

Data will be made available on request.

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.nefro.2025.501404.

References

- Suresh V, Stillman IE, Campbell KN, Meliambro K. Focal segmental glomerulosclerosis. Adv Kidney Dis Health. 2024;31:275–89.
- Rosenberg AZ, Kopp JB. Focal segmental glomerulosclerosis. Clin J Am Soc Nephrol. 2017;12:502–17.
- D'Agati VD, Fogo AB, Bruijn JA, Jennette JC. Pathologic classification of focal segmental glomerulosclerosis: a working proposal. Am J Kidney Dis. 2004;43:368–82.
- Bonilla M, Efe O, Selvaskandan H, Lerma EV, Wiegley N. A review of focal segmental glomerulosclerosis classification with a focus on genetic associations. Kidney Med. 2024;6:100826.
- Hattori M. Anti-nephrin autoantibodies: novel predictors of post-transplant recurrence of focal segmental glomerular sclerosis. Kidney Int. 2024;106:570–2.
- D'Agati VD, Kaskel FJ, Falk RJ. Focal segmental glomerulosclerosis. N Engl J Med. 2011;365:2398–411.
- Rood IM, Deegens JKJ, Wetzels JFM. Genetic causes of focal segmental glomerulosclerosis: implications for clinical practice. Nephrol Dial Transplant. 2012;27:882–90.
- 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;37: e15119
- 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 Afr Med J. 2003:445
- 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;123:123–8.
- 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.
- Ko GJ, Rhee CM, Kalantar-Zadeh K, Joshi S. The effects of high-protein diets on kidney health and longevity. J Am Soc Nephrol. 2020;31:1667–79.
- 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;322:1899.
- Bakis H, Chauveau P, Combe C, Pfirmann P. Mediterranean diet for cardiovascular risk reduction in chronic kidney disease. Adv Kidney Dis Health. 2023;30:496– 501
- Yau K, Kuah R, Cherney DZI, Lam TKT. Obesity and the kidney: mechanistic links and therapeutic advances. Nat Rev Endocrinol. 2024;20:321–35.
- McGrogan A, Franssen CFM, de Vries CS. The incidence of primary glomerulonephritis worldwide: a systematic review of the literature. Nephrol Dial Transplant. 2011;26:414–30.
- Jegatheesan D, Nath K, Reyaldeen R, Sivasuthan G, John GT, Francis L, et al. Epidemiology of biopsy-proven glomerulonephritis in Queensland adults. Nephrology. 2016;21:28–34.
- 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;12:795–800.

- 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;13e0206637.
- 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. Nephrol Dial Transplant. 1997;12:418–26.
- Molnár A, Thomas MJ, Fintha A, Kardos M, Dobi D, Tislér A, et al. Kidney biopsybased epidemiologic analysis shows growing biopsy rate among the elderly. Sci Rep. 2021;11:24479.
- 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;13:14771.
- 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 Med J. 2009;127:140–4.
- 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 J Pediatrics. 2018;14:127–33.
- Ali A, Ali MU, Akhtar SZ. Histological pattern of paediatric renal diseases in northern Pakistan. J Pak Med Assoc. 2011;61:653–8.
- Sadaf A, Khemchand M, Fouzia L, Asia Z. Clinicopathological profile of pediatric renal biopsies at a tertiary care hospital, Pakistan. Saudi J Kidney Dis Transplant. 2018;29:1403.
- Yadav S, Kandalkar B. Epidemiology of pediatric renal diseases and its histopathological spectrum – a single-center experience from India. Saudi J Kidney Dis Transplant. 2021;32:1744.
- 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;3:100061.
- 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;66:920–3.
- 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;127:1715–20.
- Zhou Fd, Zhao Mh, Zou Wz, Liu G, Wang H. The changing spectrum of primary glomerular diseases within 15 years: a survey of 3331 patients in a single Chinese centre. Nephrol Dial Transplant. 2008;24:870–6.
- 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;38:1021–30.
- 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. 2018:40:657–64.
- 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. 2022:17e0272237.
- 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 Dis. 2018;4:10–9.
- 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. Clin J Am Soc Nephrol. 2018;13:1047–54.
- 37. Su S, Yu J, Wang Y, Wang Y, Li J, Xu Z. Clinicopathologic correlations of renal biopsy findings from northeast China. Medicine. 2019;98:e15880.
- Wang Y, Zhang L, Yuan L, Xie Q, Liu S, ming HC. Changes in the spectrum of biopsy-proven renal diseases over 11 years: a single-center study in China. Ren Fail. 2024;46:2381614, http://dx.doi.org/10.1080/0886022X. 2024.2381614.
- 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;2019:1–9.
- 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;1:57-62.
- 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;10:10994.
- **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;47:546–54.
- 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). 2003:116:1377–81.
- Xiong M, Wang L, Liu X, Yue S, Dong J, Li Y, et al. Kidney biopsies in elderly Chinese patients: a nationwide survey. Am J Kidney Dis. 2020;76:295–7.
- 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;81066-1066.
- 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;21:250.
- 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;26:199.
- 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;25:55–62.

- 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 J Kidney Dis Transplant. 2021;32:364.
- Kanodia KV, Vanikar AV, 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;7:e25473, http://dx.doi.org/10.5812/numonthly.25473.
- Gopaliah L, Sudakaran I, Nalumakkal S, Narayanan R, Vareed B. Spectrum of biopsy-proven renal diseases: a single center experience. Saudi J Kidney Dis Transplant. 2018;29:392.
- 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;23:168.
- Zahir Z, Wani A, Jain M, Agrawal V, Jain S. Pediatric glomerular diseases in North India – epidemiology and clinicopathologic correlation. Indian J Nephrol. 2023;33:28.
- Bhawane A, Pasari AS, Tolani P, Balwani MR. Spectrum of biopsy-proven native kidney disease in central India. Saudi J Kidney Dis Transplant. 2022;33:688–92.
- 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;15:493–503.
- 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;16:903–20.
- 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;19:496–505.
- 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;11:2602.
- 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;36:2257–67.
- 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;17:155–73.
- 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;26:199.
- 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;37:510–4, http://dx.doi.org/10.12669/ nims.37.2.3535.
- 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;42:247.
- 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. Nephrol Dial Transplant. 2009;24:2406–10.
- 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;39:540–6.
- 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;39:60–9.
- 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;18:202–9.
- 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 J Kidney Dis Transplant. 2018:29:658.
- Pakfetrat M, Malekmakan L, Torabinezhad S, Yousefi O, Naddaffard D. Review of renal biopsies, a single center experience. Iran J Kidney Dis. 2020;14:12–9.
- 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 J Kidney Dis Transplant. 2023;34:346–54.
- 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.
- Abdullah LS. Histopathological pattern of pediatric renal diseases: a study from a university hospital in western Saudi Arabia. Saudi J Kidney Dis Transpl. 2012;23:377–84.
- 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 J Kidney Dis Transplant. 2013;24:1265.
- 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;41:933–9.
- 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;38:1228–33.
- 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;21:481.

- Almardini R, Albaramki J, Al-Saliata G, Farah M, AlRabadi K, Albderat J. Pediatric focal segmental glomerulosclerosis in Jordan: a tertiary hospital experience. Saudi J Kidney Dis Transplant. 2018;29:816.
- Sheyyab A, Al-thnaibat M, Zghayer AA, Alsheyyab J, Hamed R. Common glomerular diseases in adult Jordanians: a single-center experience. Int J Nephrol. 2022:2022:1–5.
- 79. Subedi M, Bartaula B, Pant AshokR, Adhikari P, Sharma Sanjib K. Pattern of glomerular disease and clinicopathological correlation: a single-center study from Eastern Nepal. Saudi J Kidney Dis Transplant. 2018;29:1410.
- Garyal, Kafle RK. Hisopathological spectrum of glomerular disease in Nepal: a seven-year retrospective study. Nepal Med Coll J. 2008;10:126–8.
- 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. J Nepal Med Assoc. 2023;61:163–6.
- 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. Nephrol Dial Transplant. 2010;25:3962–9.
- 83. Aoun M, Halabi C, Ammar W. Treatment of glomerular diseases in Lebanon. Saudi J Kidney Dis Transplant. 2021;32:1089.
- 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;11:148– 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;19:6.
- 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;89 Suppl. 2:S106–11.
- Alwahaibi N, Al-Khazimi O, Al-Riyami M. Histopathological spectrum of glomerular diseases in Oman: a five-year study. Saudi J Kidney Dis Transplant. 2022;33:425–31.
- 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;19:257.
- Habib M, Badruddoza S. Pattern of glomerular diseases among adults in Rajshahi, the Northern Region of Bangladesh. Saudi J Kidney Dis Transplant. 2012;23:876.
- 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. 2023:24:181
- AlYousef A, AlSahow A, AlHelal B, Alqallaf A, Abdallah E, Abdellatif M, et al. Glomerulonephritis histopathological pattern change. BMC Nephrol. 2020;21:196
- 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 Dis. 2019;5:247–58.
- Stratta P, Segoloni GP, Canavese C, Sandri L, Mazzucco G, Roccatello D, et al. Incidence of biopsy-proven primary glomerulonephritis in an Italian Province. Am J Kidney Dis. 1996;27:631–9.
- 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;20:329–35.
- 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;66:890–4.
- Zaza G, Bernich P, Lupo A. Incidence of primary glomerulonephritis in a large North-Eastern Italian area: a 13-year renal biopsy study. Nephrol Dial Transplant. 2013;28:367–72.
- 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. Nephrol Dial Transplant. 2006;21:419–24.
- 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;14:148.
- 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;54:365–76.
- 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;35:254–8.
- 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. Nephrol Dial Transplant. 2017;32 Suppl. 2:ii209–18.
- Rivera F. Frequency of renal pathology in Spain 1994–1999. Nephrol Dial Transplant. 2002;17:1594–602.
- 103. López-Gómez JM, Rivera F. Registro de glomerulonefritis de la Sociedad Española de Nefrología en 2019: pasado, presente y nuevos retos. Nefrología. 2020;40:371– 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. Front Nephrol. 2023;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. Nephrol Dial Transplant. 2004;19:3040–9.

- 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;28:39–49.
- Heaf J, Løkkegaard H, Larsen S. The epidemiology and prognosis of glomerulonephritis in Denmark 1985–1997. Nephrol Dial Transplant. 1999;14:1889–97.
- 108. Heaf JG, Sørensen SS, Hansen A. Increased incidence and improved prognosis of glomerulonephritis: a national 30-year study. Clin Kidney J. 2021;14:1594–602.
- 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. Nephrol Dial Transplant. 2008;24:877–85.
- 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;25:1054–9.
- 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. 2013:45:1577–87.
- 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;15:1361–72.
- 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 Nephrol. 2023;38:1533–45.
- 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;46:1192–8.
- 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;43:1117–26.
- Oygar DD, Neild GH. Reporting renal biopsies from Cyprus: a systematic approach. J Nephropathol. 2017;6:231–9.
- 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;47:655–62.
- 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;46:389–94.
- 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. Braz J Nephrol. 2024;46. e20230143.
- Wirta O, Mustonen J, Helin H, Pasternack A. Incidence of biopsy-proven glomerulonephritis. Nephrol Dial Transplant. 2007;23:193–200.
- 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 Nephrol. 2023;38:1249–56.
- 122. Hanko JB, Mullan RN, O'Rourke DM, McNamee PT, Maxwell AP, Courtney AE. The changing pattern of adult primary glomerular disease. Nephrol Dial Transplant. 2009;24:3050–4.
- 123. Swaminathan S, Leung N, Lager DJ, Melton LJ, Bergstralh EJ, Rohlinger A, et al. Changing incidence of glomerular disease in Olmsted County, Minnesota. Clin J Am Soc Nephrol. 2006;1:483–7.
- 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;92:1772–81.
- 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. Am J Kidney Dis. 2016;68:533–44.
- 126. Nair R, Walker PD. Is IgA nephropathy the commonest primary glomerulopathy among young adults in the USA? Kidney Int. 2006;69:1455–8.
- Murugapandian S, Mansour I, Hudeeb M, Hamed K, Hammode E, Bijin B, et al. Epidemiology of glomerular disease in Southern Arizona. Medicine. 2016;95: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;65:441–5.

- 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. Nephrol Dial Transplant. 2010;25:490–6.
- 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. J Bras Nefrol. 2017:39:29–35.
- 131. Malafronte P, Mastroianni-Kirsztajn G, Betonico GN, Romao JE, Alves MAR, Carvalho MF, et al. Paulista registry of glomerulonephritis: 5-year data report. Nephrol Dial Transplant. 2006;21:3098–105.
- 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;22:414.
- 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;22:414.
- Arias LF, Jiménez CA, Arroyave MJ. Histologic variants of primary focal segmental glomerulosclerosis: presentation and outcome. J Bras Nefrol. 2013;35:112–9.
- 135. Barrera-Herrera LE, López Panqueva RdP, 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. Rev Esp Patol. 2017;50:3–7.
- 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;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 J Kidney Dis Transplant. 2022;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;150:403–8.
- 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;52054358118799690, http://dx.doi.org/10.1177/2054358118799690.
- 140. Valjalo R, Mallea MT. Caracterización de enfermedades glomerulares: análisis de 22 años de biopsias renales. Rev Med Chil. 2023;151:52–60.
- 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;25:113–20.
- 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;53:325–32.
- 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;16:213.
- 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;26:16.
- 145. Ibrahim S, Fadda S, Fayed A, Belal D. A five-year analysis of the incidence of glomerulonephritis at Cairo University Hospital-Egypt. Saudi J Kidney Dis Transplant. 2012;23:866.
- 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. Nephrol Dial Transplant. 2011;26:1853–61.
- 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;22:257.
- 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;21:778–83.
- 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;82:1321–31.
- 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;28:32–5.