

Increasing Incidence of Inadequate Kidney Biopsy Samples Over Time: A 16-Year Retrospective Analysis From a Large National Renal Biopsy Laboratory



Caleb J. Nissen^{1,2,4}, Vanessa Moreno^{1,4}, Vicki G. Davis³ and Patrick D. Walker¹

¹Arkana Laboratories, Little Rock, Arizona, USA; ²University of Arkansas for Medical Sciences, Little Rock, Arkansas, USA; and ³Research Affiliate, Arkana Laboratories, Little Rock, Arkansas, USA

Introduction: Renal biopsy remains an essential tool for the diagnosis and treatment of patients with medical kidney disease. Recently, there has been a perceived change in the number of inadequate samples. The aim of this study was to determine the native renal biopsy miss rate from 2005 to 2020 at Arkana Laboratories, a nationwide kidney biopsy service.

Methods: From 2005 to 2020, a total of 123,372 native kidney biopsies were received from >2500 nephrologists practicing across 44 US states. The miss rate was determined by age and year. In a subset of biopsies received in 2005 and 2018, the biopsy operator was determined, nephrologist or radiologist. Furthermore, the miss rate, needle gauge, biopsy depth by operator, and biopsy core width by gauge were measured.

Results: The miss rate increased markedly from 2% in 2005 to 14% in 2020. Radiologists performed 5% of biopsies in 2005 and 95% in 2018 using smaller diameter (18g/20g) needles 92% of the time. Glomeruli per centimeter of core biopsy and mean core width were significantly lower with smaller needles. The miss rate deep was significantly lower for nephrologists and remained consistent within operator between the 2 time points. The miss rate did not correlate with the increasing age of the population who had biopsies.

Conclusion: This increase in kidney biopsy miss rate significantly affects patient care in the management of medical kidney disease. Its correlation with the complete reversal in operators suggests an urgent need for interaction with and training of radiologists in this critical technique.

Kidney Int Rep (2022) 7, 251–258; <https://doi.org/10.1016/j.ekir.2021.11.026>

KEYWORDS: interventional nephrology; interventional radiology; medical renal disease; percutaneous renal biopsy
© 2021 International Society of Nephrology. Published by Elsevier Inc. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Although the first detailed study of the percutaneous renal biopsy (PRB) was published in a Cuban journal in 1950, the publications by Iversen and Brun and the modification by Kark and Muehrcke led to its widespread use.^{1–3} By 1960, the rapid advances in the understanding of renal pathology brought on by PRB, along with significant progress in renal dialysis, transplantation, and the substantially increased understanding of renal physiology, led to the formation of Nephrology as a separate discipline.⁴ Today, PRB remains a critical tool for accurate diagnosis and thus treatment of medical renal disease.⁵

As in all biopsy procedures, safety and yield remain the critical outcome determinants. In the last 16 years,

we retrospectively observed a decline in tissue adequacy. By 2015, there were certain centers with insufficient material in as high as 50% of biopsies performed. Analysis of these centers revealed a change in operators from nephrologists to radiologists. The time frame correlates with the shift in operators nationwide.^{6,7}

The purpose of this study was to evaluate the change in renal biopsy yield in the last 16 years in our laboratory. Our hypothesis is that there has been a significant decrease in biopsy tissue obtained and that this correlates with the change to radiologists as operators and the attendant use of smaller gauge needles.⁸

Correspondence: Patrick D. Walker, Arkana Laboratories, 10810 Executive Center Drive, Street 100, Little Rock, Arkansas 72211, USA. E-mail: patrick.walker@arkanalabs.com

⁴CJN and VM contributed equally to this work.

Received 12 November 2021; accepted 22 November 2021; published online 8 December 2021

METHODS

Institutional Board Review

The Solutions Institutional Review Board approved this study as minimal risk research as the data collected were those typically obtained for routine clinical

practice. Thus, the requirement for informed consent was waived.

Histology

All samples were processed using standard techniques as in our previous studies.^{9,10} Biopsy samples for light microscopy were fixed and transported in neutral-buffered formalin. The tissue was dehydrated in a step-wise fashion in graded alcohol solutions. The alcohol was removed with graded xylene solutions, and the tissue was embedded in paraffin. Serial 3 μ sections were cut and stained with hematoxylin and eosin, periodic acid–Schiff, Jones methenamine silver, or Masson's trichrome using standard reagents.

Miss Rate Over Time

All native renal biopsies received at Arkana Laboratories from January 1, 2005 to December 31, 2020, were analyzed to determine the number of inadequate samples on light microscopy. Biopsies were considered inadequate when the diagnosis line included any of the following terms: "inadequate for diagnosis," "insufficient for diagnosis," "medulla only," "no kidney tissue available," "fat," and/or "connective tissue" and/or "skeletal muscle" only and/or ≤ 7 glomeruli.¹¹ Cases with a diagnosis including the terms "limited sample" or "limited tissue" or "limited material" were evaluated for adequacy on a case-by-case basis by one of us (PDW).

Miss Rate by Operator

In 2005, the operator was known in 93% of the cases. During that year, Arkana (then Nephropath) received biopsies from 338 nephrologists from 24 states. Nevertheless, the number of biopsies received in 2006 was 41% greater than in 2005 and the operator could only be determined in 42% of the cases. Analysis of 200 consecutive cases received in mid-2008 found that the operator could only be determined in 36% of the cases. Thus, an accurate analysis of misses by operator could not be performed after 2005. To address misses by operator at a later time point, native kidney biopsies received in a 20-week period during April to August of 2018 were evaluated. The operator was determined by requesting that information from the sending facility by phone. Of the 5201 biopsies received, the operator could still not be determined in 67 cases (1%) resulting in 5134 biopsies for analysis.

The miss rate by operator and age of patient (by decade) were also determined.

Needle Gauge by Operator

For the first time point (2005), 100 consecutive native biopsies were examined for gauge and operator. In 3 cases, one or the other criterion could not be

determined leaving 97 biopsies reported (87 performed by nephrologists and 10 by radiologists). In 2018, needle gauge data were available for 20 of 660 biopsies (3%) performed by nephrologists and 374 of 12,819 biopsies (3%) done by radiologists.

Glomeruli by Needle Gauge

Glomeruli, percent cortex, and total core length were determined in 250 consecutive biopsies from 2018 using 16g or 18g needles. Of the 250 cases, 25 had $< 40\%$ cortex and were discarded leaving 225 for inclusion. Because 14g and 20g needles were much less often used, 25 consecutive biopsies using 14g needles and 20 consecutive biopsies using 20g needles with $\geq 60\%$ cortex were evaluated.

Needle Gauge Width

The biopsy width of the tissue was calculated using the serially sectioned, periodic acid–Schiff-stained slide in the middle of the slide set. There were 102 consecutive biopsies that were evaluated for gauge, width, and glomerular number for 14g, 16g, or 18g needle. Because 20g needle biopsies were less frequent, 15 consecutive cases using a 20g needle were evaluated. Mean biopsy width was determined using 5 measurements per sample with an Olympus UC90 digital camera and Olympus cellSens image analysis software on an Olympus BX51 microscope with 100 \times objective (Olympus Corporation, Tokyo, Japan).

Miss Depth

The microscopic description in the report was used to determine the nature of the tissue received (reviewed by PDW). A deep miss was defined as medulla. A shallow miss was defined as perirenal tissue, such as muscle, fat, and/or connective tissue. In both settings, scant renal cortex was occasionally present.

Statistical Analysis

Descriptive statistics are presented with counts, percentages, and a bar chart for categorical variables, means, and SEM and box plots for continuous variables. A one-way analysis of variance test was used to evaluate the number of glomeruli per centimeter obtained from different needle gauges, controlling for all pairwise comparisons using the Tukey-Kramer adjustment for multiplicity. The Cochran-Armitage statistic was used to test for a trend over time in the biopsy miss rate. A Pearson correlation with 95% CI was used to describe the relationship between the width of the renal biopsy core and the needle gauge used to obtain it. The Cochran-Mantel-Haenszel test was used to test for differences in needle gauge choice and deep misses between operators, controlling for year of biopsy.

RESULTS

Biopsy Demographics

Arkana Laboratories received 1749 PRBs from 338 referring nephrologists across 24 US States in 2005 (Figure 1). By 2012, there were 5528 PRBs from 1252 nephrologists in 32 US States. In 2020, the last year of the study, there were 14,210 native PRBs from 2466 nephrologists in 44 US states. Taken together, 123,372 biopsies were analyzed from 2005 to 2020.

Miss Rate Over Time

The miss rate in 2005 was 2% of biopsies received, whereas in 2020, it was 14%, with a notable increase in the 2009 miss rate of 9% compared with the 3% miss rate in 2008 (Figure 2). The overall miss rate for the 16-year time frame was 11%, lowest in 2005 to 2007 at 2% and highest in 2013, 2014, and 2020 at 14%. The trend test is highly significant ($Z = -26.20$, $P < 0.001$), indicating the miss rate has generally been increasing in the last 20 years.

Change in Operator Over Time

Nephrologists performed almost all medical renal biopsies received by Arkana Laboratories in the early 2000s accounting for 95% in 2005 with a miss rate of 1%. Radiologists did only 82 PRBs in 2005 but had 14 misses (17%). In 2018, radiologists performed 95% of PRBs with a miss rate of 13% whereas nephrologists, performing 5% of PRBs, missed 8% of the time (Table 1). The miss rate was lower in each group proportional to the number of biopsies performed. Nevertheless, comparing the miss rate by nephrologists

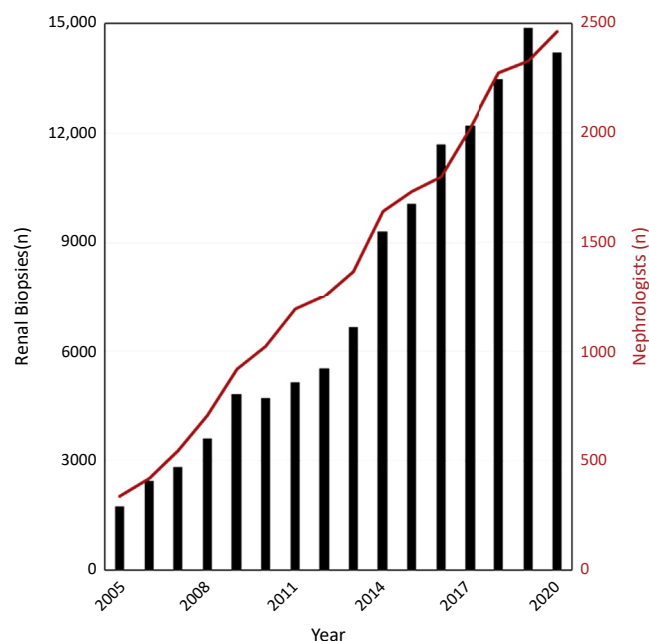


Figure 1. Native renal biopsies (bars) referred by individual nephrologists (line) across the United States over time.

Miss Rate Over Time

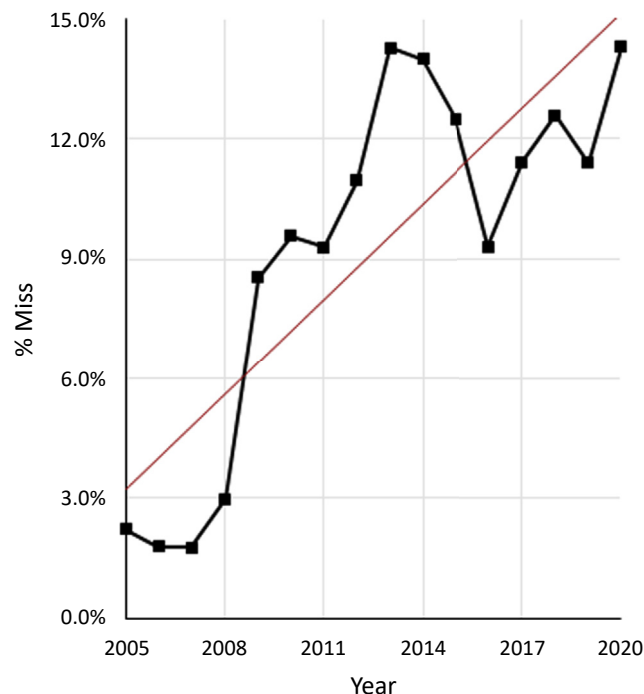


Figure 2. Miss rate over time determined by report review with the trend line in red (Cochran-Armitage trend test: $Z = -26.20$, $P < 0.001$).

when they were the primary operators (2005, 1%) with radiologists as the primary operators (2018, 13%), the overall miss rate has increased by >800% (Table 1).

Miss Rate by Operator and Age of Patient Over Time

There was a significant increase in biopsies performed on patients aged ≥ 60 years over time (Figure 3). The Cochran-Armitage trend test for 2005 to 2020 comparing $>60+$ versus <60 was significant ($Z=35.49$, $P < 0.001$). Similarly, a direct comparison of the age distributions in 2005 and 2018 also revealed an older population in 2018 (Cochran-Mantel-Haenszel $\chi^2(1) = 82.50$, $P < 0.001$), primarily driven by approximately a 10% increase in the native biopsies of 60- to 79-year-olds. Comparing miss rate by operator and age of patient in 2005, there is no difference in miss rate for either group in the older age group compared with their mean miss rate. In 2018, both nephrologists and radiologists miss more frequently in the older age group than their mean miss rate (Table 2). Nevertheless, when the miss rates by the predominate operator are compared (nephrologists in 2005 and radiologists in 2018), the miss rate for patients aged ≥ 60 years old is comparable with the overall miss rate.

Conversely, we observed a decrease in pediatric (ages 0–19) biopsies over time (Figure 3). In 2008, 14% of biopsies were from this group, whereas in 2020,

Table 1. PRB miss rate by operator and overall

Yr	PRB miss rate		
	Nephrologist <i>n/N</i> (%)	Radiologist <i>n/N</i> (%)	Overall <i>n/N</i> (%)
2005	25/1667 (1)	14/82 (17)	39/1749 (2)
2018	50/660 (8)	1648/12,819 (13)	1698/13,479 (13)

PRB, percutaneous renal biopsy of native kidneys.
Red font denotes miss rate for the primary operators that year.

pediatric biopsies accounted for only 4% of all biopsies. The Cochran-Armitage trend test comparing age groups <20 versus 20+ was significant ($Z = 28.50$, $P < 0.001$) for the years 2005 to 2020. In 2005, the miss rate was 1% for pediatric patients (1 of 145) and 2% for adult patients (38 of 1603). This rate increased for both sets of patients by 2018: 7% for pediatric patients (18 of 244) and 13% for adult patients (630 of 4890). The miss rate for adult patients was twice that of the pediatric patients for both years.

Needle Gauge by Operator

In general, nephrologists use larger needles (smaller gauge numbers) than radiologists (Table 3, $\chi^2(1) = 195$, $P < 0.001$). In 2005, nephrologists were the primary operators and used mainly 14g (22%) and 16g needles (76%). In 2018, radiologists were the primary operators and mostly preferred the 18g (86%) and 20g needles (7%).

Glomeruli per Centimeter by Gauge

A total of $N = 270$ samples were analyzed to evaluate needle gauge impact on the number of glomeruli observed per centimeter of core biopsy. Box-and-whisker plots are presented revealing the mean

(denoted by X), median (50%), interquartile range (25% and 75%), and minimum/maximum; there were no outliers detected in these data (Figure 4).

The mean number of glomeruli per centimeter is inversely related to needle gauge size, dropping from $25 \pm 0.9/cm$ to $2 \pm 1.0/cm$ (mean \pm SEM) as the gauge size increases from 14g to 20g. The pairwise comparisons between each set of gauges are significantly different based on a one-way analysis of variance with the Tukey-Kramer correction applied for multiple comparisons (all $P < 0.01$).

Needle Gauge Width

The width of the renal biopsy core changes dramatically as the needle bore size decreases (Figure 5a, Pearson's $r = -0.91$, 95% CI $[-0.94$ to $-0.87]$). The mean width of the renal tissue obtained with differing needle gauges was determined (Table 3): 14g— 894 ± 20.0 , 16g— 563 ± 10.1 , 18g— 303 ± 29.0 , and 20g— 155 ± 22.5 (mean \pm SEM in μm). A 14g biopsy sample fills the photographic field at 100 \times , and the available tissue rapidly declines with smaller bore needles (Figure 5b). Given that the mean width of a glomerulus from a healthy adult is approximately 250 μm , the bore of the most often used 18g needle is only approximately 1.2 \times larger than a glomerulus.

Biopsy Depth

In 2005, nephrologists performed 1667 biopsies (95%) missing 25 times (1%), 14 (56%) of which were deep. Radiologists did 82 biopsies (5%) and missed 14 times (17%). Of these 14, 11 (79%) were deep. In 2018, radiologists performed 12,819 (95%) and had 1648 (13%)

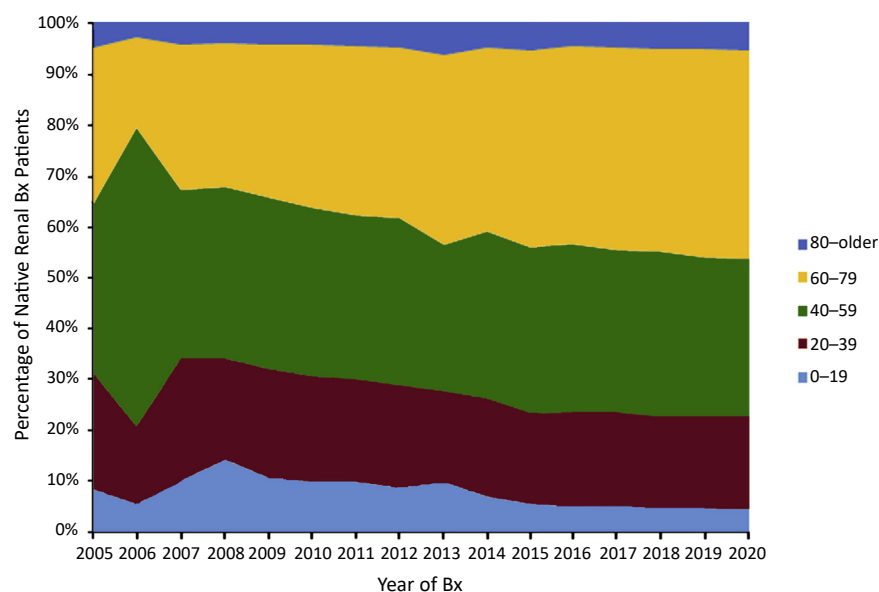


Figure 3. Native renal biopsies by age over time. The data are expressed as percentage biopsies by age group per year. There is a significant increase over time of patients aged >60 years. Comparing 2005 and 2018, there is an older population driven primarily by the almost 10% increase in patients aged 60 to 79 years (Cochran-Mantel-Haenszel $\chi^2(1) = 82.50$, $P < 0.001$). Bx, biopsy.

Table 2. PRB miss rate by operator and patient age, 2005 vs. 2018

	2005, %	2018, %
Radiologists		
Age 0–19	0	7
Age 20–39	16	13
Age 40–59	20	12
Age ≥60	17	14
Total	17	13
	2005, %	2018, %
Nephrologists		
Age 0–19	0.7	7
Age 20–39	1.9	5
Age 40–59	1.8	7
Age ≥60	1.2	10
Total	1.5	8
	Nephrologists	Radiologists
	2005, %	2018
Predominate Operator		
Age 0–19	0.7	7
Age 20–39	1.9	13
Age 40–59	1.8	12
Age ≥60	1.2	14
Total	1.5	13

PRB, percutaneous renal biopsy of native kidneys.

Red font highlights the miss rate in patients aged ≥60 years; blue font highlights miss rate across all ages.

misses, 1340 (81%) deep. Nephrologists did 660 biopsies (5%) with 50 (8%) misses, 29 (58%) deep in that same year. The 23% higher rate of deep misses for radiologists compared with nephrologists, controlling for year, is highly significant ($\chi^2(1) = 18.33, P < 0.001$).

DISCUSSION

The greatest strength of this study is its size and breadth of data. More than 120,000 native renal biopsies referred by >2500 nephrologists from 44 US States were evaluated in the 16-year study period. On this basis, these data are likely the best indicator of PRB miss rate in the United States. Though this breadth markedly increases the likelihood that this is representative of the overall renal biopsy miss rate, it does conceal the individual operator and operator group outcomes. Looking at these subsets, the miss rate varies from <1% to >50% (data not shown). It also conceals

improvement over time owing to training and consistent review of outcomes. Several groups have improved from >40% miss rate to <2% miss rate, 18g and 20g needles to all 16g needles, and a mean of <10 glomeruli/cm to >30 glom/cm with a marked decrease in significant complications (data not shown). The cause(s) of the variability in miss rate after the initial 3-fold increase in 2009 could not be determined. Still, the overall increase from a 2% miss rate (2005) to a 14% miss rate (2020) is a 7-fold increase and represents thousands more patients with insufficient tissue when, ideally, one miss is too many.

This increased miss rate correlates with the change from nephrologists (95%, 2005) to radiologists (95%, 2018) as primary operators. The lack of operator data in the entire study is a weakness. Nevertheless, this is significantly mitigated by the data that were gathered on a case-by-case basis from 2005 to 2018, the published national trends regarding the switch to radiologists as operators^{7,12–14} and the extensive anecdotal analysis of our database regarding miss rates in centers that have switched from primarily nephrologists to entirely radiologists as operators (data not shown). The introduction of the automated biopsy gun and improved visualization techniques led to a radical change in ease and safety of the PRB. With radiologists controlling access to ultrasound and scanning instruments and because the biopsy gun was so much faster, easier to use, and safer, more and more biopsies were done by radiologists. Still, nephrologists have practically been forced to hand off this technique owing to increased regulations, liability costs, and the time constraints imposed by using radiology's imaging facilities.^{7,12}

The change from nephrologists to radiologists as the primary operators in PRBs is associated not only with an overall increased miss rate but an increased miss rate by nephrologists in our study. This may reflect a decrease in the number of biopsies performed by an individual practicing nephrologist and/or inadequate training as this transition has taken place in training centers.^{6,12,15} The latter has been disputed by Korbet *et al.*⁷ in their review of the change from nephrologists to radiologists as operators doing PRBs in their center

Table 3. Biopsy core diameter and needle gauge use by operator and year

Needle gauge	Biopsy core diameter (μm , mean \pm SEM)	Nephrologist		Radiologist	
		2005 ($n = 87$) n (%)	2018 ($n = 20$) n (%)	2005 ($n = 10$) n (%)	2018 ($n = 374$) n (%)
14g	894 \pm 20.0	19 (22)	4 (20)	0 (0)	1 (0.3)
16g	563 \pm 10.1	66 (76)	16 (80)	5 (50)	24 (6)
18g	303 \pm 29.0	2 (2)	0 (0)	5 (50)	322 (86)
20g	155 \pm 22.5	0 (0)	0 (0)	0 (0)	27 (7)

Cochran-Mantel-Haenszel test for differences in operator's choice of needle gauge, controlling for year: $\chi^2(1) = 195, P < 0.001$. Red font denotes needle gauge usage for the primary operators that year.

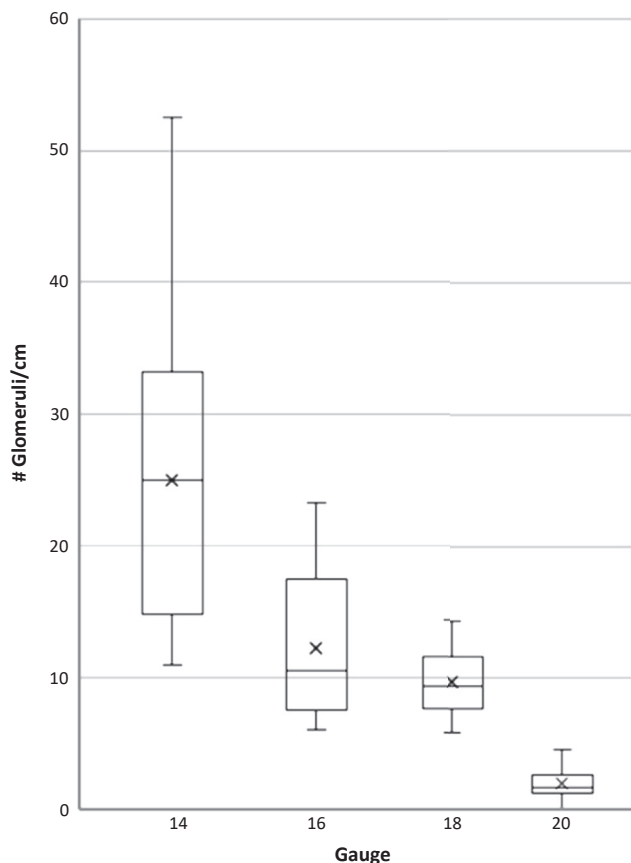


Figure 4. Number of glomeruli/cm biopsy core by needle gauge. The data are expressed as the median (middle line) in a box bounded by the IQR 25%–75%, the mean (x), and the min and max lines. 14g—med 25.0, mean 25.1, IQR 14.8–33.2, min/max 11.0–52.5, n=25; 16g—med 10.6, mean 12.3, IQR 7.6–17.5, min/max 6.2–23.3, n=40; 18g—med 9.4, mean 9.8, IQR 7.7–11.7, min/max 5.9–14.4, n=185; 20g—med 1.8, mean 2.1, IQR 1.3–2.7, min/max 0.1–4.6, n=20. In pairwise comparisons from a one-way analysis of variance, the mean number of glomeruli/cm from each needle gauge is significantly different from all other means; all $P < 0.01$ controlling for multiple comparisons. #, number; IQR, interquartile range; max, maximum; min, minimum.

and the review of the Walter Reed training program outcomes regarding competency in the performance of the PRB by Yuan *et al.*¹²

Another explanation for the marked increase in biopsy miss rate is the possibility that biopsies were done in older patients with more chronic diseases. These patients would have a thinner cortex increasing the likelihood of a miss unrelated to operator. We document that there has been a significant increase in biopsies performed on patients aged ≥ 60 years. As previously discussed, in 2018, nephrologists miss significantly more often when the patient is 60 years or older compared with their overall miss rate. Nevertheless, whether comparing the nephrologists, the radiologists, or most operators (nephrologists in 2005 and radiologists in 2018), the miss rate for patients aged ≥ 60 years is very similar to the overall miss rate

in both years. Further confirmation that a decrease in cortical thickness owing to age and/or chronic kidney disease is not related to the increased miss rate is the 10-fold increase in the miss rate among pediatric patients between nephrologists as primary operators in 2005 and radiologists as primary operators in 2018.

In our study, the needle biopsy gauge changed significantly between 2005 and 2018. The 18g needle is now the most common size by far and even 20g needles are used. The increased use of smaller 18g needles correlates with the change from nephrologists to radiologists as the primary operators both in our study and in other reports.^{6,7,12,16} The rationale for using a smaller gauge is likely based on an intuitive but incorrect assumption that smaller means safer. In a recent systematic review and meta-analysis of 87 manuscripts describing $>118,000$ PRBs, Poggio *et al.*¹⁷ found a numerical trend toward more hematomas and transfusions with 18g needles and a significant increase in pain with the 18g needles when compared with 16g needles. Nevertheless, there are also multiple studies revealing that the safety of the 14g needle is not less than either 16g or 18g needles.^{18–20} The 20g needle, while useful in endoscopic biopsies of the liver and pancreas, has a core diameter less than the diameter of a glomerulus and produces significantly less volume of tissue that is much more fragile and fragmented. Therefore, the 20g needle should not be used for a PRB.^{5,21–23}

Confirming previous studies, our data reveal that the 18g needle produces fewer glomeruli/cm than the larger core needles.^{16,20,24} Although significantly different, the number of glomeruli/cm obtained is comparable (mean \pm SEM: 18g— 9.8 ± 0.32 ; 16g— 12.3 ± 0.69). This similarity may be related to the wide variability in the bevel of needles collectively referred to only by gauge number.^{22,25} Nevertheless, there are 3 other important factors that greatly reduce the use of the smaller 18g needle of any bevel. The smaller needle produces greater fragmentation of the sample impairing an accurate evaluation of the tubulointerstitial compartment, the most important area for patient prognosis.²⁶ It has also been found that as many as 50% of glomeruli are lost or floating in 18g biopsies.²⁷ Finally, the smaller volume of an 18g sample is such that fewer total sections can be obtained. This is revealed by the significantly lower mean width of the renal core obtained by 18g and 20g needles compared with 14g and 16g needles. This lack of tissue can be critical in focal segmental lesions (focal segmental glomerulosclerosis, lupus nephritis, vasculitis, crescentic glomerulonephritis, etc.) that will more often be missed and because deeper sections for additional special stains are more often unavailable.²⁸

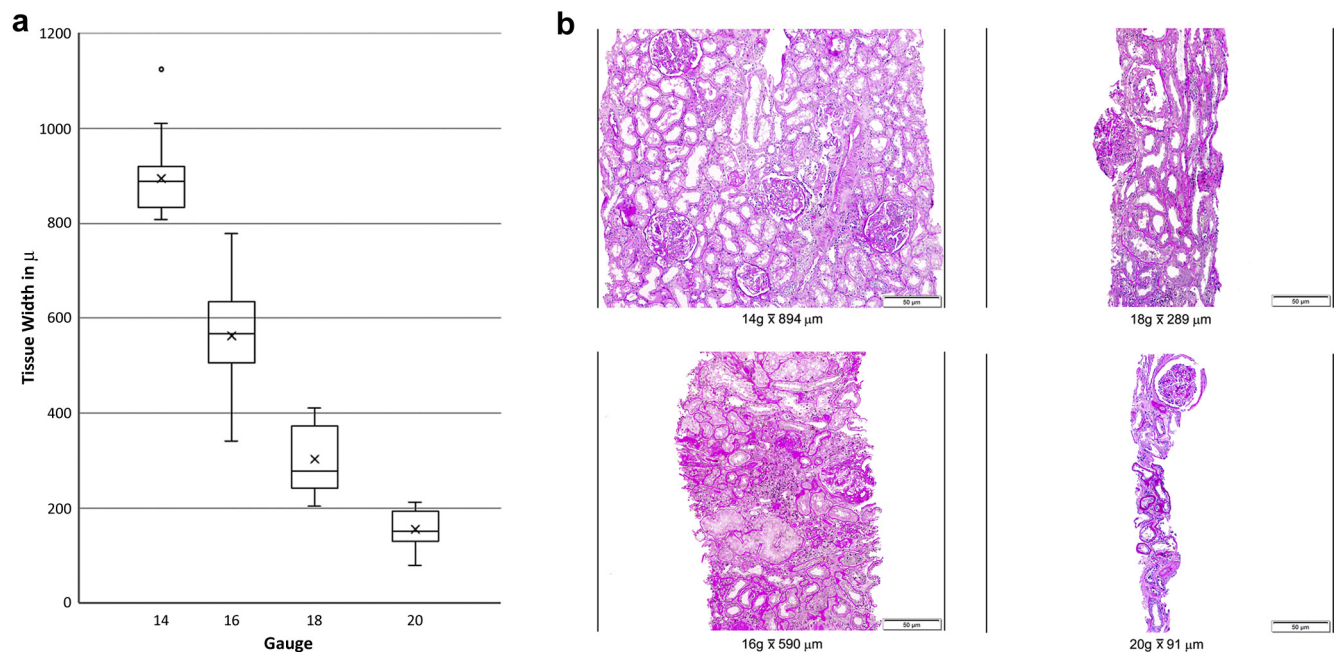


Figure 5. Width of renal core by needle gauge. (a) Box plot of tissue width in microns. Box bounded by first and third quartiles, center bar = median, x = mean, whiskers = minimum and maximum values, circle = outlier (mean \pm SEM: 14g—894 \pm 20.0, 16g—563 \pm 10.1, 18g—303 \pm 29.0, 20g—155 \pm 22.5). Pearson's correlation between needle gauge and tissue width: $r = -0.91$, 95% CI [-0.94 to -0.87]. (b) Representative photomicrographs of renal cores obtained with different needle gauges revealing example relative widths: 14g—888 μm , 16g—565 μm , 18g—325 μm , and 20g—174 μm . Note that the 20g core is less than the mean width of the glomerulus. Tissue compression followed by decompression during the procedure allowed the single glomerulus to be obtained (original magnification $\times 100$; the bars on either side of each micrograph represent the photographic field of view).

Biopsy depth plays a role in the avoidance of significant bleeding complications. A cutting needle of any gauge that passes through a medium or large artery is likely to cause serious bleeding.^{20,29} Given that the medulla contains larger vessels and is very rarely the location of the diagnostic material, it is considered inadequate for diagnosis and potentially leads to serious bleeding. In our study, the miss rate deep is markedly higher when a radiologist is the operator (>80% of misses). Nevertheless, nephrologists miss deep almost 60% of the time. As a result of Arkana being an independent laboratory, we are unable to obtain information regarding complications from the PRB; this is a major weakness that cannot be remedied.

The ability to make a diagnosis on limited tissue was not evaluated as this does not relieve the operator of the requirement to provide an adequate sample. Rarely, the diagnosis can be made on 1 glomerulus (e.g., membranous glomerulopathy). Nevertheless, even when this occurs, the biopsy remains an inadequate sample in that a possible second diagnosis, the degree of global sclerosis, tubulointerstitial fibrosis, and arterial disease all remain indeterminate.

This is the largest ever study of renal biopsy adequacy for diagnosis from a single laboratory receiving samples from a highly representative sample of nephrologists and radiologists across the United States. In

it, we document the markedly increased miss rate over time correlated with the change in operator from almost entirely nephrologists to almost entirely radiologists. This change also correlates with an increased miss rate deep and the use of the smaller 18g needle that has been found to result in less glomeruli/cm kidney core and less overall renal volume.

Given that it is unlikely that this change in operator will be reversed, one solution lies in outreach to radiologists to inform them of this marked increase in the miss rate of cortical material required for the diagnosis of medical renal disease. Then, in combination with the nephrologists, radiologists, and renal pathologists, to offer intensive short course training in correct technique. That such training can improve outcome has been documented anecdotally among radiology groups that have received hands-on training by one of us (PDW) or by departmental intensive internal training and follow-up of biopsy quality.³⁰ A regional workshop format has already proven successful (<http://kidneycon.org/>). Though this event focuses on nephrologists and nephrology trainees, radiologists and radiology residents would benefit equally from such a short, hands-on session devoted to the performance of kidney biopsy. Regardless of how it is done, a rapid intervention is required to reverse the significantly increased miss rate of the medical renal biopsy.

DISCLOSURE

All the authors declared no competing interests.

REFERENCES

- Pérez-Ara A. La biopsia puntural del riñón no megálico: Consideraciones generales y aportación de un nuevo método. *Bol Liga Contra Cancer*. 1950;25:121–147.
- Iversen P, Brun C. Aspiration biopsy of the kidney. *Am J Med*. 1951;11:324–330. [https://doi.org/10.1016/0002-9343\(51\)90169-6](https://doi.org/10.1016/0002-9343(51)90169-6)
- Kark RM, Muehrcke RC. Biopsy of kidney in prone position. *Lancet*. 1954;1:1047–1049. [https://doi.org/10.1016/s0140-6736\(54\)91618-9](https://doi.org/10.1016/s0140-6736(54)91618-9)
- Cameron JS, Hicks MJ. The introduction of renal biopsy into nephrology from 1901 to 1961: a paradigm of the forming of nephrology by technology. *Am J Nephrol*. 1997;17:347–358. <https://doi.org/10.1159/000169122>
- Luciano RL, Moeckel GW. Update on the native kidney biopsy: core curriculum 2019. *Am J Kidney Dis*. 2019;73:404–415. <https://doi.org/10.1053/j.ajkd.2018.10.011>
- Korbet SM. Nephrology and the percutaneous renal biopsy: a procedure in jeopardy of being lost along the way. *Clin J Am Soc Nephrol*. 2012;7:1545–1547. <https://doi.org/10.2215/CJN.08290812>
- Korbet SM, Whittier WL, Rodby RA. Changing trends in the performance of percutaneous renal biopsy from nephrologist to interventional radiologist: a single-center experience. *Am J Nephrol*. 2018;48:326–329. <https://doi.org/10.1159/000493925>
- Whittier WL, Korbet SM. Who should perform the percutaneous renal biopsy: a nephrologist or radiologist? *Semin Dial*. 2014;27:243–245. <https://doi.org/10.1111/sdi.12215>
- Larsen CP, Boils CL, Cossey LN, Sharma SG, Walker PD. Clinicopathologic features of membranous-like glomerulopathy with masked IgG kappa deposits. *Kidney Int Rep*. 2016;1:299–305. <https://doi.org/10.1016/j.ekir.2016.08.012>
- Walker PD, Cavallo T, Bonsib SM. Ad Hoc Committee on Renal Biopsy Guidelines of the Renal Pathology Society. Practice guidelines for the renal biopsy. *Mod Pathol*. 2004;17:1555–1563. <https://doi.org/10.1038/modpathol.3800239>
- Jeong HJ. Diagnosis of renal transplant rejection: Banff classification and beyond. *Kidney Res Clin Pract*. 2020;39:17–31. <https://doi.org/10.23876/j.krcp.20.003>
- Yuan CM, Nee R, Little DJ, et al. Survey of kidney biopsy clinical practice and training in the United States. *Clin J Am Soc Nephrol*. 2018;13:718–725. <https://doi.org/10.2215/CJN.13471217>
- Tondel C, Vikse BE, Bostad L, Svarstad E. Safety and complications of percutaneous kidney biopsies in 715 children and 8573 adults in Norway 1988–2010. *Clin J Am Soc Nephrol*. 2012;7:1591–1597. <https://doi.org/10.2215/CJN.02150212>
- Gupta RK, Balogun RA. Native renal biopsies: complications and glomerular yield between radiologists and nephrologists. *J Nephrol*. 2005;18:553–558.
- Berns JS. Training nephrology fellows in temporary hemodialysis catheter placement and kidney biopsies is needed and should be required. *Clin J Am Soc Nephrol*. 2018;13:1099–1101. <https://doi.org/10.2215/CJN.00040118>
- Sousanieh G, Whittier WL, Rodby RA, Peev V, Korbet SM. Percutaneous renal biopsy using an 18-gauge automated needle is not optimal. *Am J Nephrol*. 2020;51:982–987. <https://doi.org/10.1159/000512902>
- Poggio ED, McClelland RL, Blank KN, et al. Systematic review and meta-analysis of native kidney biopsy complications. *Clin J Am Soc Nephrol*. 2020;15:1595–1602. <https://doi.org/10.2215/CJN.04710420>
- Franke M, Kramarczyk A, Taylan C, Maintz D, Hoppe B, Koerber F. Ultrasound-guided percutaneous renal biopsy in 295 children and adolescents: role of ultrasound and analysis of complications. *PLoS One*. 2014;9:e114737. <https://doi.org/10.1371/journal.pone.0114737>
- Korbet SM, Volpini KC, Whittier WL. Percutaneous renal biopsy of native kidneys: a single-center experience of 1,055 biopsies. *Am J Nephrol*. 2014;39:153–162. <https://doi.org/10.1159/000358334>
- Nicholson ML, Wheatley TJ, Doughman TM, et al. A prospective randomized trial of three different sizes of core-cutting needle for renal transplant biopsy. *Kidney Int*. 2000;58:390–395. <https://doi.org/10.1046/j.1523-1755.2000.00177.x>
- Armellini E, Manfrin E, Trisolini E, et al. Histologic retrieval rate of a newly designed side-bevelled 20G needle for EUS-guided tissue acquisition of solid pancreatic lesions. *U Eur Gastroenterol J*. 2019;7:96–104. <https://doi.org/10.1177/2050640618804443>
- Schulman AR, Thompson CC, Odze R, Chan WW, Ryou M. Optimizing EUS-guided liver biopsy sampling: comprehensive assessment of needle types and tissue acquisition techniques. *Gastrointest Endosc*. 2017;85:419–426. <https://doi.org/10.1016/j.gie.2016.07.065>
- Roth R, Parikh S, Makey D, et al. When size matters: diagnostic value of kidney biopsy according to the gauge of the biopsy needle. *Am J Nephrol*. 2013;37:249–254. <https://doi.org/10.1159/000347219>
- Mai J, Yong J, Dixon H, et al. Is bigger better? A retrospective analysis of native renal biopsies with 16 gauge versus 18 gauge automatic needles. *Nephrol (Carlton)*. 2013;18:525–530. <https://doi.org/10.1111/nep.12093>
- Sey MS, Al-Haddad M, Imperiale TF, McGreevy K, Lin J, DeWitt JM. EUS-guided liver biopsy for parenchymal disease: a comparison of diagnostic yield between two core biopsy needles. *Gastrointest Endosc*. 2016;83:347–352. <https://doi.org/10.1016/j.gie.2015.08.012>
- Cassol CA, Braga JR, Dabbo S, Khalili K, Avila-Casado C. Effectiveness and safety of two 18-gauge needle types on native and allograft renal biopsies. *Ann Diagn Pathol*. 2017;28:1–6. <https://doi.org/10.1016/j.anndiagpath.2017.02.002>
- Van Damme B, Van Damme-Lombaerts R, Waer M. Biopsy device for obtaining kidney specimens. *Pediatr Nephrol*. 1990;4:94–95. <https://doi.org/10.1007/BF00858450>
- Komaiko MS, Jordan SC, Querfeld U, Goodman MD. A new percutaneous renal biopsy device for pediatric patients. *Pediatr Nephrol*. 1989;3:191–193. <https://doi.org/10.1007/BF00852909>
- Beckingham IJ, Nicholson ML, Bell PR. Analysis of factors associated with complications following renal transplant needle core biopsy. *Br J Urol*. 1994;73:13–15. <https://doi.org/10.1111/j.1464-410x.1994.tb07449.x>
- Geldenhuys L, Nicholson P, Sinha N, et al. Percutaneous native renal biopsy adequacy: a successful interdepartmental quality improvement activity. *Can J Kidney Health Dis*. 2015;2:8. <https://doi.org/10.1186/s40697-015-0043-z>