Access to health care among status Aboriginal people with chronic kidney disease

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Abstract

Background: Ethnic disparities in access to health care and health outcomes are well documented. It is unclear whether similar differences exist between Aboriginal and non-Aboriginal people with chronic kidney disease in Canada. We determined whether access to care differed between status Aboriginal people (Aboriginal people registered under the federal Indian Act) and non-Aboriginal people with chronic kidney disease.

Methods: We identified 106,511 non-Aboriginal and 1182 Aboriginal patients with chronic kidney disease (estimated glomerular filtration rate less than 60 mL/min/1.73 m²). We compared outcomes, including hospital admissions, that may have been preventable with appropriate outpatient care (ambulatory-care–sensitive conditions) as well as use of specialist services, including visits to nephrologists and general internists.

Results: Aboriginal people were almost twice as likely as non-Aboriginal people to be admitted to hospital for an ambulatory-care–sensitive condition (rate ratio 1.77, 95% confidence interval [CI] 1.46–2.13). Aboriginal people with severe chronic kidney disease (estimated glomerular filtration rate < 30 mL/min/1.73 m²) were 43% less likely than non-Aboriginal people with severe chronic kidney disease to visit a nephrologist (hazard ratio 0.57, 95% CI 0.39–0.83). There was no difference in the likelihood of visiting a general internist (hazard ratio 1.00, 95% CI 0.83–1.21).

Interpretation: Increased rates of hospital admissions for ambulatory-care–sensitive conditions and a reduced likelihood of nephrology visits suggest potential inequities in care among status Aboriginal people with chronic kidney disease. The extent to which this may contribute to the higher rate of kidney failure in this population requires further exploration.

Une version française de ce résumé est disponible à l’adresse www.cmaj.ca/cgi/content/full/179/10/1007/DC1

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Ethnic disparities in access to health care are well documented; however, the majority of studies include black and Hispanic populations in the United States. The poorer health status and increased mortality among Aboriginal populations than among non-Aboriginal populations than among non-Aboriginal popula-
Methods

Study population and data sources

We used computerized laboratory data from 6 of the 9 geographically defined health regions in Alberta, Canada, to identify the study cohort. Over 80% of the province’s population lives in these 6 regions. We included residents of Alberta aged 20 and older who had 1 or more outpatient measurement of their serum creatinine level made during a 1-year period from July 1, 2003, to June 30, 2004. We excluded patients with a clinically implausible serum creatinine measurement (< 25 µmol/L). Because we were interested in stable chronic kidney disease and to avoid including episodes of acute renal failure, we also excluded laboratory measurements associated with a hospital admission. The date of the first serum creatinine measurement was used as the index date.

The study cohort was linked to provincial administrative health data to identify status Aboriginal people as well as obtain details regarding health care resource use. The term “status Aboriginal” refers to any individual registered under the federal Indian Act. Status Aboriginal people are identifiable within the Alberta Health Care Insurance Plan Registry. The registry was searched from Apr. 1, 1993, to Mar. 31, 2005, and any individual with a status Aboriginal indicator at any time was classified as “status Aboriginal” (hereinafter referred to as Aboriginal). All other people were classified as non-Aboriginal. Aboriginal people who are not registered under federal Indian Act (e.g., unregistered Aboriginal and Metis people) were included in the non-Aboriginal group. According to the 2001 census, about 70% of the Aboriginal population in Alberta is status Aboriginal.

We excluded patients who had received a kidney transplant (identified from Provincial Renal Program databases) and those receiving long-term dialysis (identified from administrative data) before their index date. Patients with diabetes mellitus were identified by use of a validated administrative algorithm. We estimated socio-economic status using the neighbourhood income-per-person equivalent, an estimate of household income adjusted for household size based on data provided by the 2001 Canadian census. We obtained location of residence, based on community size, from census data. Rural residence was defined as living in a community of less than 10 000 people.

This study was approved by the institutional review board at the University of Calgary.

Measure of kidney function

We estimated glomerular filtration rate using the abbreviated prediction equation from the Modification of Diet in Renal Disease study. This equation includes variables for age, sex, black ethnic background (v. white) and serum creatinine measurement. Although ethnic background was not available from the data sources, less than 1% of the Alberta population is black. Therefore, the impact at the population level of eliminating ethnic background from the equation was expected to be minimal. Preliminary studies have validated the use of this equation in the Aboriginal population and in a community-based population with chronic kidney disease. We standardized the serum creatinine measurements made at

| Characteristic                      | Aboriginal† (n = 1 182) | Non-Aboriginal (n = 106 511) | p value‡ |
|------------------------------------|-------------------------|------------------------------|----------|
| Age, yr, median (IQR)              | 60 (50–70)              | 71 (60–80)                   | < 0.001  |
| Female                             | 743 (63.0)              | 67 741 (63.6)                | 0.61     |
| Estimated GFR, median, mL/min/1.73 m² (IQR) | 50 (38–56)              | 52 (44–57)                   | < 0.001  |
| Estimated GFR, mL/min/1.73 m²      |                         |                              |          |
| 30–59                              | 1 014 (85.8)            | 99 594 (93.5)                |          |
| 15–29                              | 105 (8.9)               | 5 907 (5.6)                  |          |
| < 15                               | 63 (5.4)                | 1 010 (1.0)                  |          |
| Diabetes mellitus                  | 500 (42.3)              | 19 929 (18.7)                | < 0.001  |
| Rural residence                    | 550 (46.5)              | 14 208 (13.3)                | < 0.001  |
| Median household income            |                         |                              |          |
| 1st quintile (lowest)              | 566 (53.7)              | 20 613 (20.1)                |          |
| 2nd quintile                       | 198 (18.8)              | 20 948 (20.5)                |          |
| 3rd quintile                       | 143 (13.6)              | 21 231 (20.7)                |          |
| 4th quintile                       | 73 (7.0)                | 18 931 (18.5)                |          |
| 5th quintile (highest)             | 74 (7.0)                | 20 634 (20.2)                |          |

Note: GFR = glomerular filtration rate, IQR = interquartile range.
*Unless stated otherwise.
†Includes people registered under the federal Indian Act (status Aboriginal). Aboriginal people who are not registered under the federal Indian Act (e.g., unregistered Aboriginal and Metis people) were included in the non-Aboriginal group.
‡Determined by use of a rank-sum test for medians and χ² test for proportions.
We evaluated access to care during the study period (date of index measurement of estimated glomerular filtration rate to Mar. 31, 2005) by use of 2 measures: admission to hospital for an ambulatory-care–sensitive condition related to chronic kidney disease, and likelihood of a nephrology visit for severe chronic kidney disease. Ambulatory-care–sensitive conditions can typically be managed effectively in an ambulatory setting; thus, admission to hospital for such a condition reflects a potentially preventable complication resulting from inadequate access to or quality of outpatient health care.6,17 We used a modified Delphi process (with 3 Delphi rounds) and an expert panel of 12 nephrologists to identify chronic kidney disease specific and relevant ambulatory-care–sensitive conditions based on the primary discharge diagnosis. Cause-specific hospital admissions were determined by use of hospital discharge coding performed by trained individuals in accordance with the International Classification of Disease, ninth revision, clinical modification (ICD-9-CM) and the International Statistical Classification of Diseases and Related Health Problems, tenth revision, Canada (ICD-10 CA) (Appendix 1, available at www.cmaj.ca/cgi/content/full/179/10/1007/DC2).

Table 2: Rates and rate ratios for admission to hospital for ambulatory-care–sensitive conditions among Aboriginal and non-Aboriginal people with chronic kidney disease

| Analysis        | Rate per 100 person-years (95% CI) | Rate ratio (95% CI) | p value |
|-----------------|------------------------------------|---------------------|---------|
|                 | Aboriginal*                        | Non-Aboriginal       |         |
| Unadjusted      | 7.7 (6.4–9.2)                      | 2.8 (2.7–2.9)        | 2.72 (2.27–2.37) | < 0.001 |
| Adjusted†       | 2.8 (2.3–3.4)                      | 1.6 (1.5–1.6)        | 1.77 (1.46–2.13) | < 0.001 |
| Adjusted‡       | 2.4 (2.0–3.0)                      | 1.5 (1.4–1.6)        | 1.58 (1.30–1.92) | < 0.001 |

Note: CI = confidence interval.
*Includes people registered under the federal Indian Act (status Aboriginal). Aboriginal people who are not registered under the federal Indian Act (e.g., unregistered Aboriginal and Metis people) were included in the non-Aboriginal group.
†Adjusted for age, sex, diabetes, baseline estimated glomerular filtration rate and admission to hospital for a non-ambulatory-care–sensitive condition.
‡Adjusted for age, sex, diabetes, baseline estimated glomerular filtration rate, median household income quintile, admission to hospital for a non-ambulatory-care–sensitive condition, and rural location of residence.

We categorized the index rate according to the Kidney Disease Outcome Quality Initiative classification (≥ 90, 60–89, 30–59, 15–29, < 15 mL/min/1.73 m²).18 Chronic kidney disease was defined as an estimated glomerular filtration rate of less than 60 mL/min/1.73 m².

Statistical analysis

We used χ² and nonparametric (rank-sum) tests to compare differences in baseline characteristics for Aboriginal and non-Aboriginal people during the study period. For hospital admission rates for ambulatory-care–sensitive conditions related to chronic kidney disease, we counted all events for patients with multiple hospital admissions. Person-time of follow-up was based on out-of-hospital time only (i.e., subtracting the number of days spent in-hospital from the total follow-up), with patient’s data censored at death, emigration from the province and end of the study period. We used a Poisson regression model to determine the association between ethnic background and risk of admission to hospital for an ambulatory-care–sensitive condition related to chronic kidney disease, after adjustment for age, sex, diabetes, baseline estimated glomerular filtration rate and hospital admissions for non-ambulatory-care–sensitive conditions. In a sensitivity analysis, we used a Cox proportional hazards model to determine the adjusted association between ethnic background and risk of first hospitalization for an ambulatory-care–sensitive condition. Cox proportional hazards models were used to determine the adjusted association between ethnic background and likelihood of an outpatient nephrologist visit or a general internist visit for patients with severe chronic kidney disease. Poisson and Cox regression models were adjusted for sex, age, baseline estimated glomerular filtration rate and diabetes. Finally, in a second analysis, we included a model that also adjusted for household income and rural location of residence. We decided a priori not to include these variables in our primary analysis because they may be considered a component of ethnic background and result in over-adjustment.29 Rate ratios (RR) and hazard ratios (HR) greater than 1 in these analyses indicate increased risk for the outcome. Assumptions for the Cox and Poisson regression models were tested and met.
Results

Study participants
In total, 676 660 patients had at least 1 outpatient serum creatinine measurement. We excluded 129 (0.02%) patients who had a serum creatinine level less than 25 μmol/L, 2139 (0.3%) who were receiving long-term dialysis, 739 (0.1%) who had received a kidney transplant before their index date and 565 960 (83.6%) who had an estimated glomerular filtration rate greater than 60 mL/min/1.73 m². Thus, our study included included 107 693 people with chronic kidney disease. Of these, 1182 (1.1%) were status Aboriginal people. The duration of follow-up was similar for the Aboriginal (median 1.37 years) and non-Aboriginal groups (median 1.39 years). Compared with non-Aboriginal patients, Aboriginal patients were younger and were more likely to live in a rural location in the lowest quintile of median household income (Table 1). Aboriginal people were also more likely than non-Aboriginal people to have diabetes mellitus and more severe kidney dysfunction.

Access to care
In total, 6.2% of Aboriginal people and 2.7% of non-Aboriginal people had at least 1 hospital admission for an ambulatory-care–sensitive condition related to chronic kidney disease (p < 0.001). The number of hospital admissions among those with at least 1 hospital admission was similar for Aboriginal people (median 1, interquartile range 1–2) and non-Aboriginal people (median 1, interquartile range 1–1). After adjustment for age, sex, baseline estimated glomerular filtration rate, non-ambulatory-care–sensitive hospital admissions and diabetes, Aboriginal people were almost twice as likely as non-Aboriginal people to have been admitted to hospital for an ambulatory-care–sensitive condition (RR 1.77, 95% confidence interval [CI] 1.46–2.13). In a model that also adjusted for median household income and rural location of residence, the rate ratio of hospital admissions for an ambulatory-care–sensitive condition was about 1.5 times higher among Aboriginal people than among non-Aboriginal people (RR 1.58, 95% CI 1.30–1.92) (Table 2) (Full model available online in Appendix 2 at www.cmaj.ca/cgi/content/full/179/10/1007/DC2.)

A similar proportion of Aboriginal (17.3%) and non-Aboriginal patients (16.3%) had at least 1 outpatient visit to a nephrologist during the study period (p = 0.75). Among these patients, the mean index estimated glomerular filtration rate was lower among Aboriginal people (mean 18.33 mL/min/1.73 m², standard deviation [SD] 7.54 mL/min/1.73 m²) than among non-Aboriginal people (mean 22.33 mL/min/1.73 m², SD 6.17 mL/min/1.73 m²).

Aboriginal people were 43% less likely than non-Aboriginal people to have visited a nephrologist (adjusted HR 0.57, 95% CI 0.39–0.83) (Table 3). Age was the most influential confounder in this analysis: every 1-year increase in age was associated with a 2% reduction in the likelihood of a nephrologist visit among Aboriginal people (HR for age 0.98, 95% CI 0.97–0.98). Tests for interaction between age, sex, estimated glomerular filtration rate, diabetes and ethnic background were not significant, suggesting that the association between ethnic background and likelihood of a nephrologist visit was not influenced by these characteristics. In a second adjusted analysis that included location of residence and median household residence, there was a nonsignificant decrease in the likelihood of a nephrologist visit among Aboriginal people compared with non-Aboriginal people (adjusted HR 0.68, 95% CI 0.45–1.04, p = 0.007) (Table 3).

During the study period a significantly greater percentage of Aboriginal patients had at least 1 outpatient visit to a general internist compared with non-Aboriginal people (75.6% vs. 55.8%, p < 0.001). However, after adjustment, there was no association between ethnic background and likelihood of a visit to a general internist for severe chronic kidney disease (Table 4).

A similar percentage of Aboriginal (96.5%) and non-Aboriginal people (95.8%) had at least 1 outpatient visit to a general practitioner during the study period (p = 0.23). A much greater percentage of Aboriginal people (72.2%) than non-Aboriginal people (44.0%) had at least 1 visit to an emergency department during the study period (p < 0.001).

Interpretation
Our results suggest that there are differences in the care received by status Aboriginal and non-Aboriginal people with chronic kidney disease. First, Aboriginal people with chronic kidney disease do access the health care system, as shown by

| Rate per 100 person years (95% CI) | Aboriginal† | Non-Aboriginal | Hazard ratio (95% CI) | p value |
|------------------------------------|-------------|----------------|-----------------------|--------|
| Unadjusted                         | 15.6 (10.9–22.5) | 15.3 (14.5–16.2) | 1.02 (0.71–1.47)      | 0.92   |
| Adjusted†                          | 7.8 (5.4–11.3)   | 13.8 (13.0–14.8) | 0.57 (0.39–0.83)      | 0.003  |
| Adjusted§                          | 9.3 (6.1–14.0)   | 13.5 (12.6–14.4) | 0.68 (0.45–1.04)      | 0.007  |

Note: CI = confidence interval.
†Includes people registered under the federal Indian Act (status Aboriginal). Aboriginal people who are not registered under the federal Indian Act (e.g., unregistered Aboriginal and Metis people) were included in the non-Aboriginal group.
‡Adjusted for age, sex, diabetes and baseline estimated glomerular filtration rate.
§Adjusted for age, sex, diabetes, baseline estimated glomerular filtration rate, median household income quintile and rural location of residence.
the use of general practitioner and emergency department services. However, Aboriginal people with chronic kidney disease were almost twice as likely as non-Aboriginal people to be admitted to hospital for an ambulatory-care–sensitive condition related to their chronic kidney disease. In addition, despite national guidelines recommending that people with severe chronic kidney disease visit a nephrologist, we found that Aboriginal people with severe chronic kidney disease were significantly less likely than non-Aboriginal people to have visited a nephrologist.

Although universal coverage for health care in Canada has alleviated insurance-related barriers to care,30 our results suggest that other barriers may exist among Aboriginal people, who are less likely than non-Aboriginal people to receive specialized care for chronic kidney disease. This is unlikely to be related to lack of access to the health care system in general, because almost all Aboriginal people in our study had at least 1 visit to a general practitioner. The probability of obtaining a serum creatinine measurement is also unlikely to have influenced these results, as we have reported that Aboriginal people have an increased likelihood of having a serum creatinine measurement compared with non-Aboriginal people.31 Decreased access to specialized medical care among Aboriginal people has been reported.4,10 Potential barriers, including distance from specialized care, require further study. Delayed referral to a nephrologist for severe chronic kidney disease is not unique to the Aboriginal population and has been reported in other ethnic groups.12,33 Given the increased mortality among patients with a late referral to a nephrologist,12,34,35 these results suggest suboptimal quality of care in the Aboriginal population and the need for interventions to reduce or eliminate these disparities.

Access to health care is difficult to evaluate, and ambulatory-care–sensitive conditions are commonly used as a measure of access and assessment of performance of the health care system.26–28 Barriers in access are complex and include patient, environmental and health system factors. Lower socioeconomic status has been associated with an increased likelihood of hospital admission for an ambulatory-care–sensitive condition in the United States19 and an increased frequency of physician visits in Canada.18 However, even after adjustment for median household income and rural location of residence, Aboriginal people were still 50% more likely than non-Aboriginal people to be admitted to hospital for an ambulatory-care–sensitive condition related to their chronic kidney disease. Controlling for socio-economic status and location of residence may result in over-adjustment because these factors are considered to be a component of ethnicity29 and may artificially reduce differences in rates hospital admissions for ambulatory-care–sensitive conditions. Thus, in our main analysis, we did not adjust for median household income and rural location of residence. We found a similar increased risk for all-cause hospital admissions among status Aboriginal and non-Aboriginal people (Appendix 3, available at www.cmaj.ca/cgi/content/full/179/10/1007/DC2), which suggests that the increased risk extends beyond ambulatory-care–sensitive conditions related to chronic kidney disease. However, with the present data, we are unable to determine the extent to which patient factors such as compliance with recommended treatments, including attendance at physician appointments, may have influenced the study results.

Although we did not have access to the details of the medical care provided to Aboriginal people with chronic kidney disease, lack of access to specialized care may result in suboptimal use of treatments that reduce the risk of progression of kidney disease,9,10 and may contribute to higher rates of end-stage renal disease in this population. This is further supported by our previous study that showed a lower prevalence of all stages of chronic kidney disease but a higher prevalence of more severe chronic kidney disease and increased mortality rates among status Aboriginal people than among non-Aboriginal people.31

### Limitations

Our study had several limitations. First, we did not directly calibrate serum creatinine measurements to measurements made at the Cleveland Clinic, where the Modification of Diet in Renal Disease equation for estimated glomerular filtration rate was derived. We did, however, implement a province-wide standardization of serum creatinine measurements that had been indirectly calibrated to the isotope dilution mass spectrometry reference standard using the new Modification of Diet in Renal Disease equation.32 Second, we were not able to identify Metis people and non-registered Aboriginal people, which may have resulted in misclassification of some individuals as non-Aboriginal.

| Table 4: Rates and hazard ratios for likelihood of an outpatient visit to a general internist among Aboriginal and non-Aboriginal people with severe chronic kidney disease* |
|-----------------------------------------------|----------------|----------------|----------------|----------------|
| Analysis | Rate per 100 person years (95% CI) | Hazard ratio (95% CI) |
|----------|----------------------------------|-------------------|
|          | Aboriginal† | Non-Aboriginal | Unadjusted | Adjusted† | Adjusted§ |
|          | 158.3 (132.6–188.8) | 85.5 (82.9–88.3) | 1.63 (1.36–1.95) | 0.55 |
|          | 87.6 (73.0–105.1) | 87.0 (84.2–89.9) | 1.00 (0.83–1.21) | 0.97 |
|          | 92.6 (77.1–111.4) | 86.8 (84.0–89.7) | 1.05 (0.88–1.27) | < 0.001 |

*Includes people registered under the federal Indian Act (status Aboriginal). Aboriginal people who are not registered under the federal Indian Act (e.g., unregistered Aboriginal and Metis people) were included in the non-Aboriginal group.
†Adjusted for age, sex, diabetes and baseline estimated glomerular filtration rate.
‡Adjusted for age, sex, diabetes, baseline estimated glomerular filtration rate, median household income quintile and rural location of residence.
Aboriginal patients as non-Aboriginal. However, given that the majority of the Aboriginal population in Alberta is registered under the federal Indian Act and given the size of the non-Aboriginal population, this potential misclassification would have had minimal impact on our results, and if there was a bias, it would have been toward the null hypothesis. Third, use of laboratory data to define the cohort limited our study to patients who had sought medical care and who had a serum creatinine measurement, which may limit the generalizability of our results. Finally, as with all observational studies, the possibility of residual confounding cannot be excluded. Although we were unable to directly adjust for patient-related factors such as compliance and distance to the nearest nephrologist, we were able to account for key clinical variables, including diabetes and baseline kidney function. The hazard ratios associated with ethnic background and likelihood of a nephrologist visit were substantial, and they are unlikely to be completely negated by adjustment for additional covariates.

Conclusion
The results of our study suggest potential inequities in care between status Aboriginal people and non-Aboriginal people with chronic kidney disease. The extent to which these inequities may contribute to the higher rates of end-stage kidney failure among the Aboriginal population requires further exploration. Interventions that target these disparities in care are needed.

This article has been peer reviewed.

Competing interests: None declared.

Contributors: All of the authors contributed to the conception and design of the study and to the interpretation of the data. Song Gao and Brenda Hemmelgarn performed the analysis and drafted the manuscript. All of the authors revised the article for important intellectual content and approved the final version submitted for publication.

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