Epidemiology of diabetes mellitus among First Nations and non-First Nations adults

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ABSTRACT

Background: First Nations people in Canada experience a disproportionate burden of type 2 diabetes mellitus. To increase our understanding of this evolving epidemic, we compared the epidemiology of diabetes between First Nations and non-First Nations adults in Saskatchewan from 1980 to 2005.

Methods: We used administrative databases to perform a population-based study of diabetes frequency, incidence and prevalence in adults by ethnic background, year, age and sex.

Results: We identified 8275 First Nations and 82 306 non-First Nations people with diabetes from 1980 to 2005. Overall, the incidence and prevalence of diabetes were more than 4 times higher among First Nations women than among non-First Nations women and more than 2.5 times higher among First Nations men than among non-First Nations men. The number of incident cases of diabetes was highest among First Nations people aged 40–49, while the number among non-First Nations people was greatest in those aged 70 or more years. The prevalence of diabetes increased over the study period from 9.5% to 20.3% among First Nations women and from 4.9% to 16.0% among First Nations men. Among non-First Nations people, the prevalence increased from 2.0% to 5.5% among women and from 2.0% to 6.2% among men. By 2005, almost 50% of First Nations women and more than 40% of First Nations men aged 60 or older had diabetes, compared with less than 25% of non-First Nations men and less than 20% of non-First Nations women aged 80 or older.

Interpretation: Diabetes disproportionately affects First Nations women during their reproductive years. This distinct ethnicity-based pattern suggests diverse underlying mechanisms that may include differences in the diabetogenic impact of gestational diabetes.

In Saskatchewan in 1937, diabetes was not detected among the 1500 First Nations people who underwent a tuberculosis survey. By 1990, almost 10% of the province’s First Nations adults had diabetes; by 2006, the proportion was over 20%, while it remained at about 6% in the general population. Although an increased prevalence of diabetes among First Nations people has also been documented in other Canadian provinces, only recently have consistent diabetes case definitions applied to health care system administrative databases been used to compare differences between large populations of First Nations and non-First Nations people.

We sought to describe the epidemiology of diabetes in Saskatchewan from 1980 to 2005. We reasoned that finding ethnicity-based differences in trends and patterns of type 2 diabetes over the longest period reported for a Canadian jurisdiction would help to clarify the underlying mechanisms behind known disparities and translate into more effective diabetes prevention and management initiatives.

Methods

Study population

The population of Saskatchewan was about one million people throughout the study period. About 99% (both First Nations and non-First Nations people) were beneficiaries of a universal health care system and were recorded in the Ministry of Health’s insurance registry. These health care beneficiaries made up the total annual populations for this study, as they have for other population-based research. We subdivided the annual populations into First Nations and non-First Nations people. First Nations people are indigenous to Canada and, for the purposes of this study, were identifiable in the Ministry of Health databases as individuals (both those who lived on- or off-reserve) registered under Section 6 of the Indian Act of Canada and who were assigned a 10-digit number in the Indian Registry. The proportion of all provincial health care beneficiaries represented by registered First Nations people grew from about 5% to 10% during the study period.
Most people categorized as “non-First Nations” are of European origin but about 5% are of mixed European and First Nations heritage (Metis). Less than 0.5% of the non-First Nations population are nonregistered First Nations people who are not in the Indian Registry.

This study was approved by the University of Saskatchewan Ethics Review Board.

We identified people with diabetes using a validated algorithm\(^{19}\) based on the National Diabetes Surveillance System case definitions.\(^{17}\) The case definition required one hospital discharge (hospital separation database), two physician service claims (medical claims database) or a physician service claim followed by a hospital discharge for diabetes (ICD-9 250 or ICD-10-CA E10–E14) within any 730 day period. We excluded diabetes records related to gestational records to ensure that gestational diabetes cases were not counted as diabetes cases.

We defined the diabetes incident year as the first calendar year in which the individual met the diabetes case definition (after two years with no diabetes diagnosis). The incident counts for 2004 and 2005 were underestimated because meeting the case definition required up to two years. For each year after the diabetes incident year, the case was counted as a prevalent case. Annual prevalence counts did not include incident cases. We could not distinguish between types 1 and 2 diabetes, but less than 2% of patients with new diabetes each year are under age 20,\(^{18}\) which is the age group with the highest incidence of type 1 diabetes.

### Statistical analysis

We obtained de-identified diabetes incident and prevalent case counts by five-year age group, sex and First Nations status for 1980–2005 from the Ministry of Health. We determined the annual crude, age- and sex-specific and age-standardized incidence and prevalence rates of diabetes for First

| Year | First Nations women | Non-First Nations women | Incidence ratio (95% CI) | First Nations men | Non-First Nations men | Incidence ratio (95% CI) |
|------|---------------------|-------------------------|--------------------------|-------------------|-----------------------|--------------------------|
| 1980 | 118                 | 1299                    | 5.05 (4.31–5.91)         | 79                | 1206                  | 2.38 (1.97–2.87)         |
| 1981 | 84                  | 1404                    | 3.40 (2.82–4.09)         | 69                | 1124                  | 2.23 (1.83–2.74)         |
| 1982 | 106                 | 1343                    | 4.70 (3.98–5.54)         | 88                | 1343                  | 2.71 (2.27–3.25)         |
| 1983 | 120                 | 1352                    | 5.05 (4.31–5.90)         | 89                | 1248                  | 2.48 (2.07–2.97)         |
| 1984 | 122                 | 1339                    | 5.66 (4.85–6.61)         | 90                | 1211                  | 2.65 (2.22–3.17)         |
| 1985 | 121                 | 1185                    | 6.44 (5.50–7.53)         | 86                | 1259                  | 3.00 (2.50–3.60)         |
| 1986 | 127                 | 1312                    | 5.61 (4.81–6.53)         | 104               | 1462                  | 3.15 (2.67–3.72)         |
| 1987 | 146                 | 1273                    | 5.85 (5.07–6.75)         | 117               | 1572                  | 3.37 (2.87–3.94)         |
| 1988 | 121                 | 1292                    | 4.32 (3.70–5.06)         | 114               | 1543                  | 3.32 (2.83–3.90)         |
| 1989 | 114                 | 1222                    | 3.81 (3.24–4.47)         | 105               | 1232                  | 2.79 (2.36–3.29)         |
| 1990 | 128                 | 1143                    | 4.83 (4.15–5.63)         | 89                | 1022                  | 2.43 (2.03–2.91)         |
| 1991 | 153                 | 1145                    | 5.12 (4.44–5.90)         | 102               | 1237                  | 2.86 (2.42–3.39)         |
| 1992 | 155                 | 1347                    | 4.95 (4.31–5.70)         | 117               | 1565                  | 2.54 (2.17–2.98)         |
| 1993 | 168                 | 1285                    | 5.20 (4.54–5.95)         | 127               | 1275                  | 2.75 (2.37–3.21)         |
| 1994 | 190                 | 1191                    | 5.09 (4.47–5.78)         | 134               | 1281                  | 2.95 (2.55–3.43)         |
| 1995 | 184                 | 1259                    | 4.56 (4.00–5.19)         | 150               | 1337                  | 2.97 (2.58–3.42)         |
| 1996 | 184                 | 1268                    | 4.25 (3.74–4.84)         | 137               | 1242                  | 2.59 (2.24–3.00)         |
| 1997 | 196                 | 1580                    | 3.51 (3.10–3.97)         | 159               | 1326                  | 2.50 (2.18–2.87)         |
| 1998 | 206                 | 1508                    | 3.97 (3.52–4.49)         | 203               | 1673                  | 2.99 (2.64–3.37)         |
| 1999 | 245                 | 1612                    | 4.28 (3.83–4.80)         | 208               | 1708                  | 2.92 (2.59–3.29)         |
| 2000 | 244                 | 1759                    | 3.98 (3.56–4.45)         | 202               | 1638                  | 2.73 (2.42–3.09)         |
| 2001 | 298                 | 1769                    | 4.00 (3.61–4.44)         | 216               | 1628                  | 2.46 (2.19–2.76)         |
| 2002 | 280                 | 1974                    | 3.33 (3.00–3.70)         | 246               | 1773                  | 2.44 (2.19–2.73)         |
| 2003 | 263                 | 1883                    | 3.39 (3.05–3.78)         | 249               | 1780                  | 2.63 (2.36–2.93)         |
| 2004* | 250                | 1956                    | 3.35 (3.00–3.74)         | 245               | 1849                  | 2.77 (2.48–3.09)         |
| 2005* | 223                | 1585                    | 3.11 (2.76–3.50)         | 203               | 1481                  | 2.72 (2.41–3.07)         |

Total  4546                   37403                   3729                  44903  

Note: CI = confidence interval.

*Case counts and incidence are underestimated for 2004 and 2005 because the case definition required up to 2 years.
Nations and non-First Nations adults (aged ≥ 20 years). For diabetes prevalence calculations, we included all Ministry of Health beneficiaries in the denominator. To calculate diabetes incidence, we first subtracted prevalent diabetes cases. Age-standardized rate calculations used the direct method standardized to the 1991 Canadian census population. We compared the rates in First Nations and non-First Nations people by calculating annual incidence and prevalence rate ratios.

Results

Of the 90 581 cases of incident diabetes from 1980 to 2005 (Table 1), 8275 were in First Nations people (45% men) and 82 306 were in non-First Nations people (55% men). Overall, the incidence (Table 1) and prevalence (Table 2) were more than 4 times higher among First Nations men than among non-First Nations men and more than 2.5 times higher among First Nations women than among non-First Nations men.

The highest diabetes rates were among First Nations women, while non-First Nations women had the lowest rates (Figure 1, Figure 2). There was a slight convergence in diabetes rates between First Nations men and women over time, which was associated with a decrease in the incidence of diabetes among women and an increase among men. However, the prevalence of diabetes remained more than 25% higher among First Nations women than among First Nations men by 2005. From 1980 to 2005, the prevalence of diabetes more than doubled among First Nations women (9.51% to 20.33%) and more than tripled among First Nations men (4.94% to 16.01%). Among non-First Nations people, the rates of change were similar, with an increase from 2.01% to 5.51% among women and from 2.01% to 6.24% among men.

Figure 3 shows the age-specific diabetes incident case counts and incidence over time. The most consistent finding was a progressive increase in the number of new cases within each age group during 1981–1985 and 2001–2005. Exceptions were older non-First Nations adults who experienced a

| Year | First Nations women | Non-First Nations women | Prevalence ratio (95% CI) | First Nations men | Non-First Nations men | Prevalence ratio (95% CI) |
|------|---------------------|-------------------------|--------------------------|-------------------|-----------------------|--------------------------|
| 1980 | 532 9.51 | 6 601 2.01 | 4.72 (4.62–4.83) | 323 4.94 | 6 933 2.01 | 2.46 (2.39–2.54) |
| 1981 | 635 11.10 | 7 496 2.25 | 4.94 (4.84–5.04) | 387 5.76 | 8 058 2.31 | 2.50 (2.43–2.57) |
| 1982 | 695 11.73 | 8 461 2.48 | 4.73 (4.64–4.82) | 449 6.41 | 9 138 2.59 | 2.48 (2.41–2.54) |
| 1983 | 781 12.73 | 9 311 2.68 | 4.75 (4.67–4.84) | 522 7.30 | 10 158 2.85 | 2.56 (2.50–2.62) |
| 1984 | 883 14.00 | 10 123 2.66 | 4.90 (4.82–4.99) | 594 8.06 | 11 168 3.09 | 2.60 (2.55–2.66) |
| 1985 | 972 14.90 | 10 903 3.02 | 4.93 (4.85–5.01) | 659 8.60 | 11 998 3.28 | 2.62 (2.56–2.67) |
| 1986 | 1 056 15.48 | 11 480 3.15 | 4.92 (4.84–5.00) | 711 9.11 | 12 695 3.45 | 2.64 (2.59–2.70) |
| 1987 | 1 151 15.81 | 12 185 3.30 | 4.80 (4.72–4.87) | 786 9.71 | 13 418 3.61 | 2.69 (2.64–2.74) |
| 1988 | 1 269 16.13 | 12 794 3.42 | 4.71 (4.64–4.78) | 883 10.38 | 14 207 3.80 | 2.73 (2.68–2.78) |
| 1989 | 1 347 15.77 | 13 462 3.57 | 4.47 (4.36–4.49) | 969 10.84 | 14 966 3.99 | 2.72 (2.67–2.76) |
| 1990 | 1 425 15.82 | 14 008 3.67 | 4.31 (4.25–4.37) | 1 043 11.11 | 15 555 4.13 | 2.69 (2.64–2.73) |
| 1991 | 1 514 16.30 | 14 465 3.78 | 4.32 (4.26–4.38) | 1 093 11.37 | 16 012 4.27 | 2.67 (2.62–2.71) |
| 1992 | 1 611 16.28 | 14 900 3.84 | 4.24 (4.19–4.30) | 1 156 11.49 | 16 471 4.35 | 2.64 (2.60–2.69) |
| 1993 | 1 726 16.71 | 15 496 3.95 | 4.23 (4.17–4.29) | 1 231 11.58 | 17 042 4.45 | 2.60 (2.56–2.64) |
| 1994 | 1 846 17.13 | 15 961 4.04 | 4.24 (4.18–4.29) | 1 313 11.97 | 17 603 4.59 | 2.61 (2.57–2.64) |
| 1995 | 1 982 17.25 | 16 303 4.09 | 4.22 (4.17–4.27) | 1 391 11.99 | 17 978 4.64 | 2.58 (2.55–2.62) |
| 1996 | 2 124 17.43 | 16 780 4.17 | 4.18 (4.13–4.24) | 1 500 12.24 | 18 468 4.71 | 2.60 (2.56–2.64) |
| 1997 | 2 248 17.63 | 17 188 4.26 | 4.14 (4.09–4.19) | 1 582 12.63 | 18 989 4.84 | 2.61 (2.57–2.65) |
| 1998 | 2 388 17.66 | 17 857 4.35 | 4.03 (3.99–4.08) | 1 694 12.74 | 19 609 4.93 | 2.58 (2.55–2.62) |
| 1999 | 2 531 17.81 | 18 481 4.46 | 3.99 (3.95–4.04) | 1 849 13.12 | 20 395 5.04 | 2.60 (2.57–2.64) |
| 2000 | 2 706 18.67 | 19 170 4.68 | 3.99 (3.94–4.03) | 1 981 13.80 | 21 227 5.33 | 2.59 (2.56–2.62) |
| 2001 | 2 869 18.77 | 19 876 4.81 | 3.90 (3.86–3.94) | 2 125 14.15 | 21 991 5.45 | 2.60 (2.57–2.63) |
| 2002 | 3 105 19.24 | 20 649 4.96 | 3.88 (3.84–3.92) | 2 280 14.63 | 23 042 5.65 | 2.59 (2.56–2.62) |
| 2003 | 3 302 19.76 | 21 623 5.20 | 3.80 (3.76–3.84) | 2 460 15.46 | 24 242 5.96 | 2.59 (2.56–2.62) |
| 2004 | 3 478 19.95 | 22 333 5.33 | 3.74 (3.70–3.78) | 2 624 15.60 | 25 139 6.08 | 2.57 (2.54–2.59) |
| 2005 | 3 659 20.33 | 23 280 5.51 | 3.69 (3.65–3.73) | 2 791 16.01 | 26 118 6.24 | 2.57 (2.54–2.59) |

Note: CI = confidence interval.
slight decline in the number of diabetes cases in the late 1980s and early 1990s. The most striking difference between First Nations and non-First Nations people was the age at which most new diabetes cases occurred. Those peaked in the 40–49 year age group for both First Nations men and women, while most new cases among non-First Nations people were in the oldest age group (≥ 70 years). There were more diabetes cases in First Nations women than men, particularly during the reproductive years, and there were more diabetes cases among non-First Nations men than women, particularly during middle age and older. Finally, although peak age-specific diabetes incidence and incident case counts both
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occurred in the oldest non-First Nations group (≥ 70), there was a 20-year difference between the peak in new diabetes cases (ages 40–49) and incidence (ages 60–69) among First Nations people.

Figure 4 shows that, with few exceptions, there was a progressive increase in the age-specific diabetes prevalence over time in all groups. The largest increases occurred after age 40 among First Nations people and after age 50 among non-First Nations people. Among First Nations adults, the prevalence of diabetes was highest between ages 60 and 75. In contrast, the prevalence was highest after age 80 among non-First Nations adults. By 2005, almost 50% of First Nations women and more than 40% of First Nations men aged 60 and older had diabetes, compared with the highest levels of less than 25% observed among non-First Nations people aged 80 or older.

Interpretation

This study describes the epidemiology of diabetes among First Nations and non-First Nations adults over the longest period reported for a Canadian jurisdiction. We found distinct differences between populations that extend beyond known disparities in the rates of diabetes. Thus, diabetes is a disease of young First Nations adults with a marked predilection for women. In contrast, diabetes is a disease of aging non-First Nations adults that is more common among men. These observations suggest fundamental differences between populations in the mechanisms underlying diabetes; this has widespread implications that are probably also relevant to other indigenous and developing populations.1

Our findings are consistent with reports that used similar methods6,9 to show higher prevalence of diabetes among First Nations people than among non-First Nations people in neighbouring provinces. This difference appears to be at least partly because of higher rates of overweight and obesity among First Nations people.3,19,20 Because of a longer study duration and the inclusion of more detailed incidence data, we have now shown that First Nations people also have markedly different trends and patterns of diabetes than non-First Nations people. Among non-First Nations people, the prevalence of diabetes was identical among men and

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Figure 3: Age-specific diabetes incident case counts and incidence by period.
women in 1980, but it was substantially higher among men by 2005. This corresponded to a divergence in the incidence of diabetes between the sexes, which is possibly related to a greater increase in BMI observed in Canadian men than women during this period.21 Among First Nations people, the prevalence of diabetes was almost twice as high among women as it was among men in the early 1980s, and a large absolute difference has persisted. Although there was considerable annual variation in the incidence (probably partly because of small numbers), the incidence was consistently higher among First Nations women. Differences in incidence between the sexes have diminished over time, however, and it is possible that the large decrease in diabetes incidence among women in the late 1980s was accentuated by the passage of Bill C-31 in 1985. This primarily reinstated young urban First Nations women to the Indian Registry and would have increased the corresponding study denominators.13

Despite very low rates of type 1 diabetes in North American Aboriginal people,18,22 most incident cases of diabetes occurred in young First Nations adults. Furthermore, the consistently higher rates of diabetes among First Nations women than among First Nations men was related to an excess burden of diabetes in women aged 20–49. What could account for this striking sex difference? One possibility is the higher rates of overweight and obesity among First Nations women.19,23 Another factor may be the high rates of gestational diabetes that were present before the significant occurrence of type 2 diabetes in northern First Nations communities;23 gestational diabetes is strongly linked to pre-pregnancy overweight and obesity.24 Because gestational diabetes is a predictor for type 2 diabetes in affected women,25 female populations with high rates of prepregnancy overweight and obesity and gestational diabetes could experience a resultant intragenerational increase in the rate of type 2 diabetes.

Gestational diabetes has also been implicated in an inter-

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**Figure 4:** Age-specific diabetes prevalence in 1980, 1990 and 2005.
generational “vicious cycle” by increasing the risk of type 2 diabetes among the offspring. This is supported by the early appearance of gestational diabetes among First Nations people and its association with increasing rates of high birth weight, a predictor of diabetes among First Nations people. A recommendation from the Fifth International Workshop on Gestational Diabetes was to clarify the intergenerational diabetogenic role of gestational diabetes. We believe that it is equally important to clarify its intragenerational impact and are currently attempting to gain insights into the relative contribution of each through the use of simulation modelling.

The contrasting demographic features of diabetes in First Nations and non-First Nations people have different implications for prevention, screening, management and allocation of health care resources. We highlight three examples. First, with respect to screening and primary prevention initiatives for First Nations people, our findings support an emphasis on children and young adults. We believe that there is sufficient evidence for both an intra- and intergenerational diabetogenic role of gestational diabetes to focus primary prevention initiatives on the time before and during the reproductive years of First Nations women. Programs designed to prevent gestational diabetes, ensure universal gestational diabetes screening, optimize management of diabetic pregnancies and provide follow-up initiatives for women who have experienced gestational diabetes have the potential to reduce the rate of type 2 diabetes in mothers and their offspring.

Second, the large difference in the age of diabetes onset between First Nations and non-First Nations people could contribute to distinct patterns of chronic complications because of differential mortality and differential exposure to the metabolic effects of diabetes. Although speculative, the duration of exposure to diabetes and its interaction with other variables (e.g., quality of diabetes management) might be an important determinant in the relative likelihood of developing specific diabetic complications such as diabetic end-stage renal disease. This should also be a priority area for future research.

Finally, the trends reported here indicate that the prevalence of diabetes among both First Nations and non-First Nations people is likely to continue increasing in the foreseeable future, particularly as the large cohort of children and teenagers that make up about half of the First Nations population enter young adulthood. In addition, an earlier “baby boom” among non-First Nations people is approaching the age during which it will also be at the highest risk of diabetes. We are now beginning a period in which two markedly different cohorts will simultaneously experience an increase in diabetes.

Strengths and limitations
The strengths of this study included its duration, use of a validated algorithm to identify diabetes cases, the use of data for total populations, and the ability to subdivide the population by ethnic background. We are not aware of systematic differences in strategies for diabetes screening or diagnostic criteria between First Nations and non-First Nations people. However, a decrease in diagnostic fasting plasma glucose was widely instituted in 1997 and was followed by an expected rise in diabetes incidence in all study groups.

Limitations of the study included an inability to identify Aboriginal people other than First Nations, reducing the true differences between First Nations and non-First Nations people. Second, identifying cases using administrative data is likely to underestimate the incidence and prevalence of diabetes. Third, we could not differentiate between type 1 and type 2 diabetes. However, less than 3% of all non-First Nations diabetes incident cases occurred among people aged 20–29, the adult group most likely to develop type 1 diabetes. Furthermore, type 1 diabetes is very uncommon among First Nations people, including children. Thus, inclusion of type 1 diabetes cases would have only marginally increased the rates of diabetes among non-First Nations people and would have likely reduced the true differences between First Nations and non-First Nations people.

Fourth, some prevalent cases of diabetes may have been misclassified as incident cases at the beginning of the study because of delayed diagnosis or limitations of the algorithm. This could have contributed to the initial decline in diabetes incidence observed in all groups. Finally, we were not able to determine the rate of diabetes by location. However, the rates are currently higher in rural areas and are consistently lower among northern compared with southern First Nations people.

Conclusion
This study shows marked differences in the epidemiology of type 2 diabetes between First Nations and non-First Nations people. Whether this is because of relative differences in the genetics of energy balance interacting with other differences in the environmental determinants of obesity and carbohydrate intolerance is still uncertain. Complicating this further is the emerging possibility that epigenetic phenomena may play a role. What is clear is that the rapid appearance of type 2 diabetes particularly among First Nations people and other indigenous and developing populations has been precipitated by environmental rather than genetic factors. Its long-term solution will require effective primary prevention initiatives that are population-based and driven by public health and community initiatives.

This article has been peer reviewed.

Competing interests: None declared.

Contributors: Roland Dyck conceived and designed the study, acquired, analyzed and interpreted the data, and drafted the article. Mary Rose Stang helped design the study, acquired the data. Nathaniel Osgood analyzed and interpreted the data. Ting Hsiang Lin and Amy Gao analyzed the data. Nathaniel Osgood, Ting Hsiang Lin, Mary Rose Stang, and Amy Gao revised the manuscript critically for important intellectual content. All authors gave final approval of the version submitted for publication.

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