

# Trends in glucose testing among individuals without diabetes in Ontario between 2010 and 2017: a population-based cohort study

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## Abstract

**Background:** Early identification of people with diabetes or prediabetes enables greater opportunities for glycemic control and management strategies to prevent related complications. To identify gaps in screening for these conditions, we examined population trends in receipt of timely glucose testing overall and in specific clinical subgroups.

**Methods:** Using linked administrative databases, we conducted a retrospective cohort study of people aged 40 years and older without diabetes at baseline. Our primary outcome was up-to-date glucose testing, defined as having received testing at least once in the 3 years before each index year from 2010 to 2017, using linked administrative databases of people residing in Ontario, Canada. We calculated rates of up-to-date testing by age group, sex, ethnicity (South Asian, Chinese, general population) and comorbidities (hypertension, hyperlipidemia, cardiovascular disease).

**Results:** Over the 8-year study period, up-to-date glucose testing rates were stable at 67% for men and 77% for women (both relative risk 1.00 per year; 95% confidence interval 1.00–1.00). Testing rates were significantly lower in men than in women (all age groups  $p < 0.001$ ) and lower in younger than older age groups (except those aged  $\geq 80$  yr). South Asian people had the highest testing rates, although among people aged 70 years or older, testing was highest in the general population ( $p < 0.001$ ). Among people with hypertension, hyperlipidemia and cardiovascular disease, annual testing rates were also stable, but only 58% overall among people with hypertension.

**Interpretation:** We found lower glucose testing rates in younger men and people with hypertension. Our findings reinforce the need for initiatives to increase awareness of glycemic testing.

Diabetes mellitus is a major cause of morbidity and mortality, accounting for 1.5 million deaths worldwide in 2012 and attributable to 11.9% of deaths in Canada in 2009.<sup>1–3</sup> The prevalence of diabetes has also been steadily increasing, with an estimated 3.4 million Canadians or 9.3% of the population affected in 2015.<sup>4–6</sup> By 2025, the prevalence is predicted to rise 44% to 5 million Canadians, or 12.1% of the population.<sup>5</sup> With considerable advances in glycemic control measures and management strategies — such as early lifestyle modifications and novel pharmacologic interventions — which have the potential to reduce morbidity and mortality, screening for diabetes is cost-effective.<sup>7–10</sup> However, despite advocacy for early diagnosis and intervention,<sup>11</sup> diabetes often goes unnoticed and appropriate interventions are delayed as a substantial proportion of individuals who ultimately receive diagnoses may be asymptomatic in the initial phases for many years.<sup>12,13</sup>

In Canada, the prevalence of undiagnosed diabetes is estimated to be 1.1%–3.1%.<sup>14</sup> Diabetes Canada and the Canadian Cardiovascular Harmonized National Guidelines Endeavour (C-CHANGE) guidelines thus recommend that all adults aged 40 years and older be screened every 3 years, and those at very high risk regardless of age (e.g., with cardiovascular disease or cardiac risk factors, and some ethnic groups) be screened every 6 to 12 months to ensure early diagnosis and initiation of appropriate interventions to reduce morbidity and mortality.<sup>15,16</sup>

**Competing interests:** See the end of the article.

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Several studies have described trends in prediabetes and diabetes incidence and prevalence in the general adult population, but few have assessed screening practices in the context of clinical practice guidelines.<sup>4,17–21</sup> To identify potential gaps in testing by age, sex or ethnicity, our objective was to examine temporal trends in the proportion of adults in Ontario, Canada, who were 40 years and older without a diabetes diagnosis and who received blood glucose testing (including glycosylated hemoglobin [HbA<sub>1c</sub>]) in the previous 3 years, making them up to date with screening recommendations.<sup>15,16</sup> We also examined trends among adults with hypertension, hyperlipidemia and cardiovascular disease.

## Methods

### Study design and population

We conducted a retrospective, population-based cohort study of the entire population of Ontario eligible for the province's publicly available health insurance plan (Ontario Health Insurance Plan [OHIP]) using multiple population-based health administrative databases. We identified residents without known diabetes between 2008 and 2017 from the Ontario Registered Persons Database (RPDB), which contains demographic information about eligibility for OHIP, including birth date, sex and residential postal code.

We examined glucose testing rates among both the overall population of adults and a subcohort of individuals with hypertension, hyperlipidemia and a history of cardiovascular disease.<sup>15,16</sup> For the overall population, we created annual study cohorts from 2010 to 2017 of individuals eligible for OHIP and aged at least 40 years for the entire 3 years before. Thus, individuals entered the cohort at a minimum age of 40 years, but evaluation of up-to-date glucose testing began at age 43 years.

Using the Diabetes Canada and C-CHANGE guidelines, we defined up-to-date testing as having received blood glucose or HbA<sub>1c</sub> testing at least once during those 3 years.<sup>15,16</sup> For the subcohorts of individuals with hypertension, hyperlipidemia and cardiovascular disease, for whom screening is recommended every 6 to 12 months, we examined annual testing between 2008 and 2017.<sup>15,16</sup>

### Data sources

We identified individuals with a diabetes diagnosis through linkage to the Ontario Diabetes Database (ODD), which is a database of all Ontario residents with physician-diagnosed diabetes, created using the Canadian Institute for Health Information Discharge Abstract Database (CIHI-DAD) of all acute care hospital admissions and the OHIP Claims database.<sup>22</sup> The OHIP database contains information from physician claims and diagnostic and community laboratory services in Ontario.<sup>23,24</sup> The negative predictive value of the ODD has been validated at 97.6%.<sup>22</sup>

Similarly, we identified individuals with hypertension from the Ontario Hypertension Database, a validated database of Ontario residents with physician-diagnosed hypertension.<sup>25</sup> We identified individuals with hyperlipidemia from the OHIP

database, using an algorithm of at least 2 physician claims with a diagnosis of hyperlipidemia within 2 years before the study year, and determined history of cardiovascular disease from the CIHI-DAD; codes for cardiovascular diseases are shown in Appendix 1 (available at [www.cmajopen.ca/content/10/3/E772/suppl/DC1](http://www.cmajopen.ca/content/10/3/E772/suppl/DC1)).<sup>26</sup>

For ethnicity, we used a validated surname algorithm to classify individuals into 3 groups, based on likely ethnicity: Chinese, South Asian and all others in the general population.<sup>27</sup> We determined rural or urban residence and neighbourhood income quintile through linkage of each individual's postal code to Statistics Canada Census data.<sup>28</sup> We used the Client Agency Program Enrolment (CAPE) database, maintained by the Ontario Ministry of Health to track patient enrolment with primary care providers, to identify individuals rostered to a primary care physician in a group practice.<sup>29</sup> We classified as virtually rostered those individuals who were not formally rostered to a physician in CAPE but received most of their primary care services from a particular physician.

### Outcomes

To identify people receiving glucose testing, we used the OHIP database, which provides health services data on all residents of the province, and the Ontario Laboratories Information System, and included HbA<sub>1c</sub>, serum or plasma glucose, and oral glucose tolerance testing. The Ontario Laboratories Information System is a province-wide, centralized repository of laboratory results from tests conducted in Ontario, which began in 2007.<sup>30</sup> Because OHIP claims do not differentiate between fasting and nonfasting blood glucose tests, we were unable to restrict to fasting tests as recommended in Canadian guidelines, although diagnoses can be made using random tests in situations of overt hyperglycemia.<sup>15</sup> Details of codes used to identify glucose testing are shown in Appendix 1.

We excluded tests performed while a person was in hospital because our focus was on testing in the primary care setting, and restricted OHIP claims to tests ordered by family physicians. We also excluded tests performed on pregnant women, as they undergo glycemic testing for specific pregnancy-related indications. All data sets were linked using unique, encoded identifiers and analyzed at ICES (formerly Institute for Clinical Evaluative Sciences).

### Statistical analysis

We calculated absolute rates of up-to-date testing for 2010 to 2017 overall and for test type by sex, 10-year age bands up to age 80 years or older, and ethnicity. We compared characteristics of individuals up to date with testing versus not for the latest (2017) cohort using means and standard deviations for age and number of physician visits, and proportions for categorical variables. To examine factors associated with being up to date with testing, we additionally performed logistic regression, adjusting for age, sex, rural residence, neighbourhood income quintile, ethnicity, hypertension, hyperlipidemia and cardiovascular disease, which have been identified in previous studies of health services access and outcomes.<sup>31,32</sup>

In post-hoc analyses, we calculated up-to-date testing rates in those who were and in those who were not rostered formally or virtually to a primary care physician as a proxy for access to a physician. Post hoc, we stratified analyses by rostering to primary care physician. We also examined annual testing rates for individuals with hypertension, hyperlipidemia and cardiovascular disease by sex and age group. We compared testing rates between groups using Poisson regression, modelling the number of individuals tested as a linear function of year and population, and the natural logarithm of the number of eligible individuals as the offset. We also ran separate models for each predefined subgroup to examine temporal trends. Because an individual's test in 1 year contributes to the outcome for the following 2 years (i.e., were nonindependent), we performed bootstrapping with 1000 samples to estimate 95% confidence intervals (CIs) and *p* values. We conducted all analyses using SAS version 9.4 (SAS Institute) and considered 2-sided *p* values < 0.05 to be significant.

### Ethics approval

The use of the data in this project is authorized under section 45 of Ontario's *Personal Health Information Protection Act* and does not require review by a research ethics board.

### Results

We identified about 5.5 million individuals in each 3-year study cohort. Mean age was approximately 57 years and 47% were men. Chinese people comprised about 6% and South Asian people about 3% of each cohort.

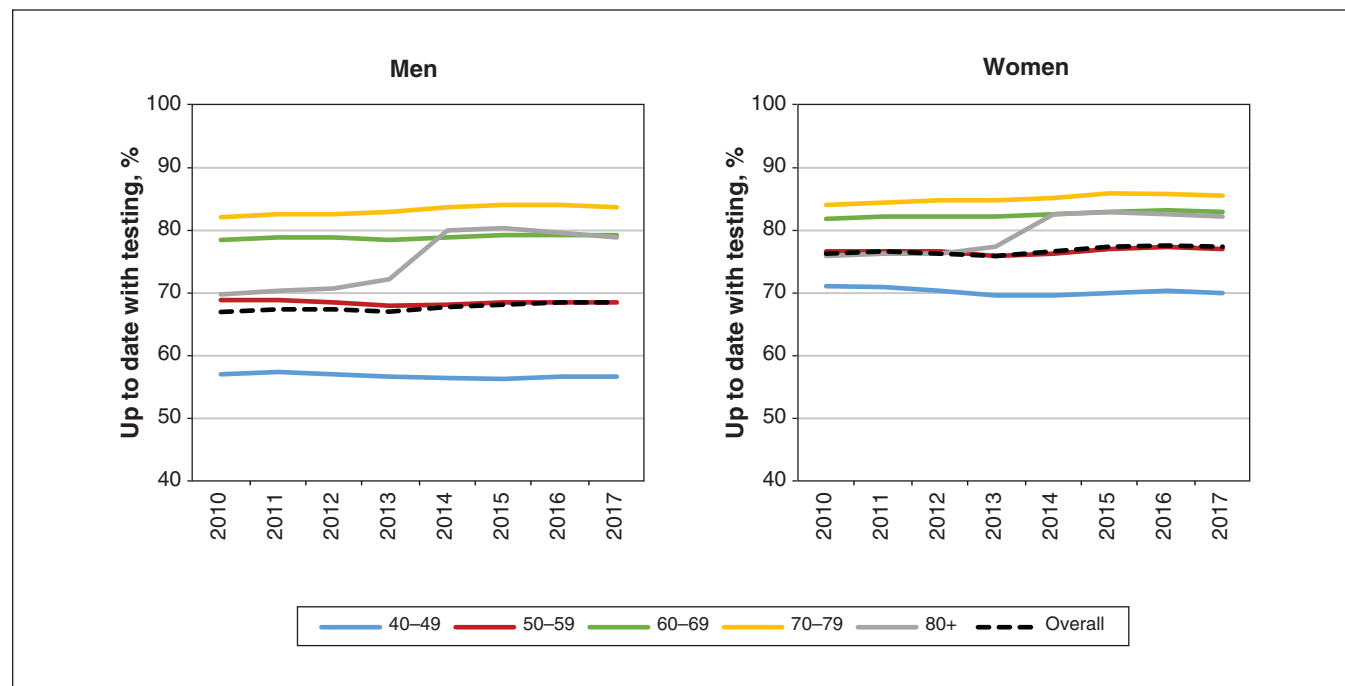
### Testing by age and sex

Figure 1 and Appendix 2 (available at [www.cmajopen.ca/content/10/3/E772/suppl/DC1](http://www.cmajopen.ca/content/10/3/E772/suppl/DC1)) show rates of up-to-date glucose testing by age group and sex overall and by test type, respectively. Over time, we observed overall glucose testing rates to be stable at 67% among men and 77% among women (both relative risk [RR] 1.00 per year; 95% CI 1.00–1.00), although we also observed increases in HbA<sub>1c</sub> testing concomitant with decreases in serum or plasma glucose testing.

Across all age groups, rates were higher among women than men (all *p* < 0.001). We observed the greatest absolute differences between men and women in the 40–49-year age group, where 57% of men were up to date with testing, compared with 70% of women. For individuals younger than 80 years, rates of up-to-date glucose testing increased with age for both men and women (combined 63% among the 40–49-year age group v. 84% among the 70–79-year age group). Among individuals aged 80 years and older, rates up to 2013 were stable at about 70% among men and 76% among women. Thereafter, rates rose sharply to relatively new stable rates of 80% and 82%, respectively, and was attributable to increases in both HbA<sub>1c</sub> and glucose testing (Appendix 2).

### Testing by ethnicity

Age- and sex-stratified results by ethnicity showed similar trends to age- and sex- stratified results in the overall population (Figure 2). A greater proportion of women were up to date with testing than men (all ethnic groups, *p* < 0.001), as were older age groups compared with younger age groups, except for Chinese and South Asian people aged 70 years or older.

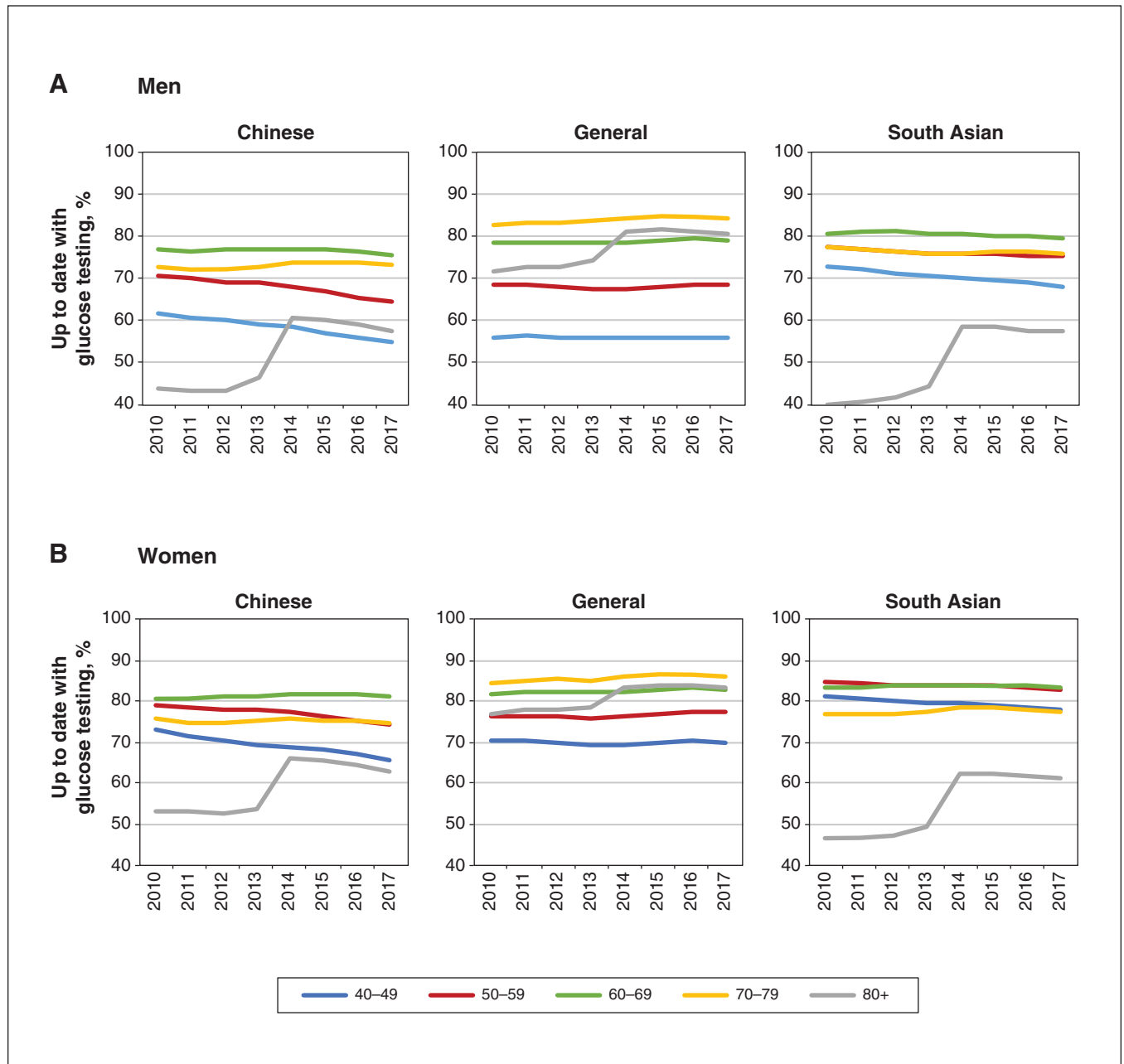


**Figure 1:** Up-to-date glucose testing rates among adults aged 40 years or older in Ontario, by age (years), 2010–2017. “Up to date” is defined as at least 1 glycosylated hemoglobin, plasma or serum glucose or oral glucose tolerance test in the previous 3 years, and excludes people who received a diabetes diagnosis before the study period.

Overall, South Asian people had the highest glucose testing rates (about 77%; RR 1.06, 95% CI 1.06–1.06, general population as reference), followed by the general (72%) and Chinese populations (about 69%; RR 0.96, 95% CI 0.95–0.96, general population as reference). However, among adults aged 70 years and older, testing was markedly higher in the general population than among the South Asian and Chinese groups ( $p < 0.001$  for both), but among men younger than 60 years, South Asian people had higher up-to-date testing rates than their Chinese and general population counterparts ( $p < 0.05$  for both).

### Up-to-date testing

Characteristics of those up to date with testing in 2017 are shown in Table 1 and Appendix 3 (available at [www.cmajopen.ca/content/10/3/E772/suppl/DC1](http://www.cmajopen.ca/content/10/3/E772/suppl/DC1)). Older age, being female, living in a high-income neighbourhood, having a regular primary care physician and visiting a physician more often were all associated with a greater likelihood of being up to date with testing ( $p < 0.001$ ). Individuals with hypertension or a history of cardiovascular disease were also more likely to have been tested in the previous 3 years than those without ( $> 85\%$ ).



**Figure 2:** Up-to-date glucose testing rates among (A) men and (B) women aged 40 years and older in Ontario, by ethnicity, 2010–2017. “Up to date” is defined as at least 1 glycosylated hemoglobin, plasma or serum glucose or oral glucose tolerance test in the previous 3 years, and excludes people who received a diabetes diagnosis before the study period. Groups are displayed in order of increasing overall rates of testing in 2017.

**Table 1: Characteristics of being up to date or not up to date with glucose testing among people aged 40 years and older in Ontario, 2017**

Characteristic	No. (%)* of population up to date with glucose testing in 2017		Std. difference†
	Yes n = 4 264 253	No n = 1 567 009	
Sex			0.22
Men	1 891 174 (44.3)	869 149 (55.5)	
Women	2 373 074 (55.7)	697 858 (44.5)	
Age, yr			
Mean ± SD	58.1 ± 12.0	53.5 ± 11.3	0.39
Median (IQR)	56 (49–66)	51 (45–59)	0.42
Community size‡			0.03
Rural	467 345 (11.0)	183 529 (11.8)	
Urban	3 787 951 (89.0)	1 370 074 (88.2)	
Neighbourhood income quintile‡			
1 (highest)	744 427 (17.5)	333 079 (21.5)	0.1
2	823 090 (19.4)	311 113 (20.0)	0.02
3	853 497 (20.1)	294 214 (18.9)	0.03
4	873 713 (20.5)	288 753 (18.6)	0.05
5 (lowest)	959 188 (22.6)	325 952 (21.0)	0.04
Surname-based ethnicity			
Chinese	233 458 (5.5)	111 486 (7.1)	0.07
South Asian	126 223 (3.0)	39 755 (2.5)	0.04
Other	3 904 567 (91.6)	1 415 701 (90.4)	0.03
Rostered to a primary care physician			
Yes	3 566 023 (83.6)	918 832 (58.6)	0.57
Virtual§	575 762 (13.5)	219 041 (14.0)	0.01
No	122 463 (2.9)	429 134 (27.4)	0.73
No. of visits to a family physician (2014–2016)			
Mean ± SD	11.7 ± 11.8	4.8 ± 8.9	0.66
Median (IQR)	9 (4–15)	2 (0–6)	1.03
No. of visits to a family physician (2014–2016): categorized			
None	200 418 (4.7)	561 869 (35.9)	0.84
1–3	626 772 (14.7)	382 643 (24.4)	0.25
4–9	1 462 081 (34.3)	386 404 (24.7)	0.21
10+	1 974 977 (46.3)	236 091 (15.1)	0.72
Hypertension	1 593 337 (37.4)	247 523 (15.8)	0.50
History of hyperlipidemia	218 957 (5.1)	13 479 (0.9)	0.25
Any history of cardiovascular disease¶	208 689 (4.9)	30 320 (1.9)	0.16

Note: IQR = interquartile range, SD = standard deviation.  
 \*Unless otherwise specified.  
 †p value for differences all < 0.001.  
 ‡Data for community size and neighbourhood income quintile are missing for 22 356 and 24 229 people, respectively.  
 §With virtual rostering, a patient is not formally on a physician's roster, but received most of their primary care from a specific physician in 2015/16.  
 ¶Defined as any previous hospital admission for myocardial infarction, stroke, heart failure, percutaneous coronary intervention or coronary artery bypass graft surgery.

## Additional analyses

We conducted post-hoc analyses to characterize our findings further. First, when we examined people rostered to a primary care physician separately from those who were not, those who were rostered were more likely to be up to date with testing (about 77% v. about 21%; Appendix 4, available at [www.cmajopen.ca/content/10/3/E772/suppl/DC1](http://www.cmajopen.ca/content/10/3/E772/suppl/DC1)). However, we observed mild increases among nonrostered, older age groups (particularly people aged 80 years and older) between 2012 and 2017.

In analyses of important clinical subgroups, annual testing rates among individuals with hypertension, hyperlipidemia and cardiovascular disease were stable during the study period (Figure 3; about 58%, 70% and 60%, respectively). Testing rates were higher among older than younger adults, and among women than men, especially among younger age groups. Among those with hypertension, about 65% of men and women aged 70–79 years were tested annually compared with 44% and 51% of men and women aged 40–49 years, respectively. Among men and women aged 70–79 years with cardiovascular disease, about 66% were tested annually compared with less than 50% of men and women aged 40–49 years; and among those with hyperlipidemia, annual testing rates were almost 80% among adults aged 70 years and older compared with 61% and 67% of men and women aged 40–49 years, respectively.

## Interpretation

In this study of contemporary trends in rates of up-to-date glucose testing over the last decade, we found that the proportion of people up to date as recommended by Canadian practice guidelines<sup>15,16</sup> has been stable among both men and women, across different age and ethnic groups, and among subpopulations of individuals with hypertension and hyperlipidemia. We identified important gaps in recommended testing among young men and older Chinese and South Asian people. Annual testing rates were also suboptimal among individuals with hypertension, hyperlipidemia and cardiovascular disease.

Notable among those aged 80 years and older was the increase in up-to-date testing rates between 2013 and 2014, possibly attributable to Diabetes Canada's 2013 release of a new guideline encouraging use of HbA<sub>1c</sub> testing.<sup>33</sup> Although screening in this population is of questionable benefit and recommended on an individual basis, without the need for prior fasting, testing became more convenient for this population, which may not tolerate fasting.<sup>34</sup> In addition, in this older population, much focus may be on the management of existing chronic conditions and comorbidities. As such, primary preventive care such as diabetes screening may have been overlooked until the guideline's release brought greater attention to the importance of glycemic testing, particularly among people without a regular primary care physician.

Our results are consistent with earlier studies of glucose testing and diabetes screening.<sup>35,36</sup> Lee and colleagues' cross-sectional study of electronic medical records found that fewer than 50% of rostered patients received screening over a 36-month period, and Wilson and colleagues reported that 63% of Ontarians aged 40 years and older without diabetes

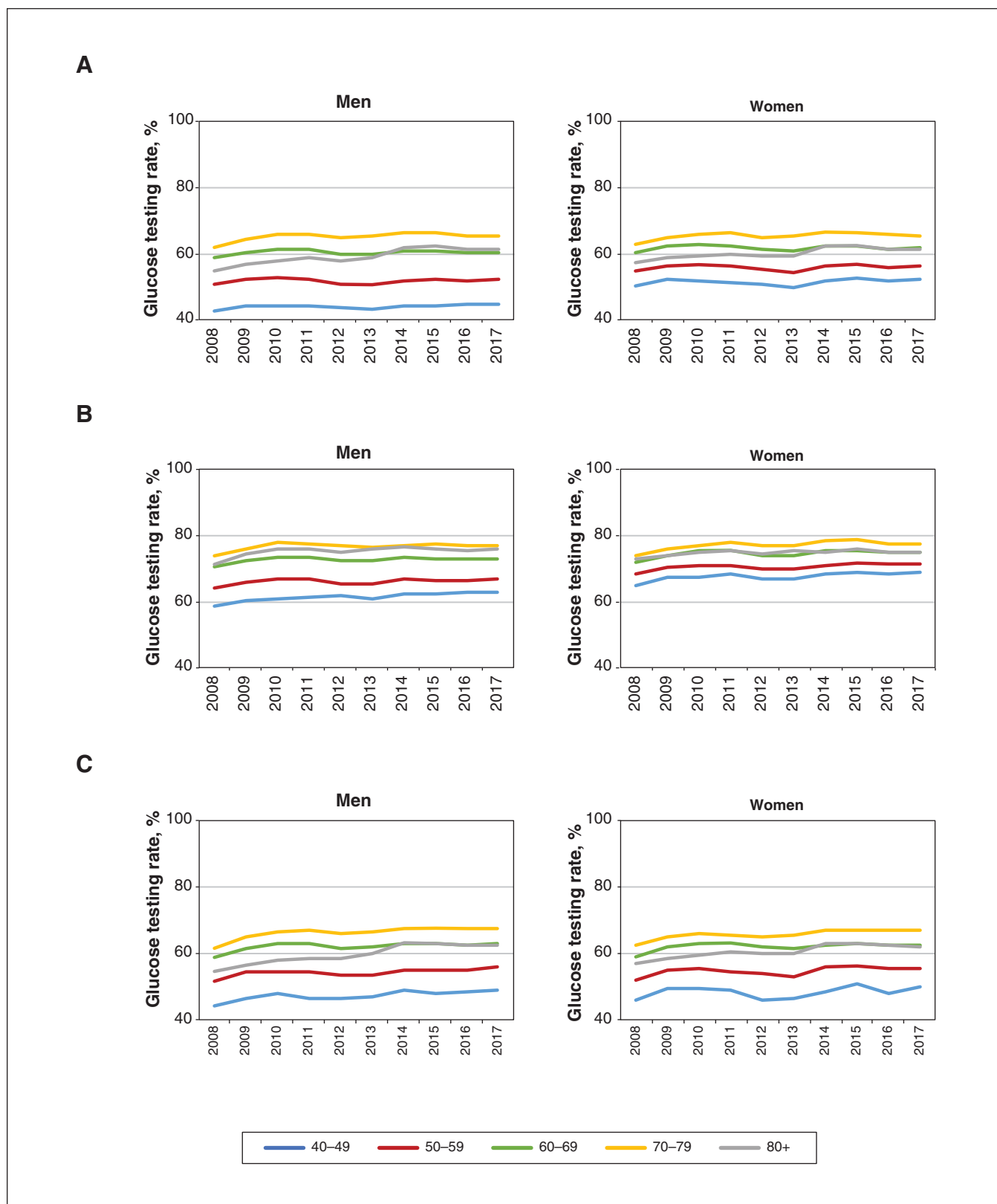
received blood glucose testing between 2003 and 2005.<sup>35–37</sup> Findings of increasing uptake of HbA<sub>1c</sub> testing between 1995 and 2015 may contribute to our higher overall testing rates.<sup>36,38</sup>

Regarding ethnicity, higher testing rates have been reported among immigrants in Ontario than among the general population, with South Asian immigrants having the highest rates.<sup>35</sup> Similar to our findings, older immigrants were also less likely to be tested than their general population counterparts. Whether this is a result of generation gaps in health-seeking behaviour, language barriers, health literacy or other reasons is unknown.<sup>39</sup> In the United States, results from studies of ethnic groups have been mixed, with some reporting lower testing rates among several visible-minority groups compared with non-Hispanic white people,<sup>40,41</sup> and another study finding no significant association between “high-risk” ethnicity and diabetes screening.<sup>42</sup> Although differences between Canada and the US may be attributable to greater access through Canada's publicly available health care system, further understanding of contributing factors to lower testing among older immigrants is required.

Studies have indicated that diabetes prevalence is on the rise globally, partly owing to increased survival.<sup>4–6</sup> However, a recent systematic review suggests that diabetes incidence, which increased in many populations up to the mid-2000s, has since stabilized or declined in many regions.<sup>43,44</sup> In Ontario, diabetes incidence increased 31% between 1997 and 2003,<sup>18</sup> yet data from the Canadian Chronic Disease Surveillance System show a decrease in incidence between 2006/07 and 2013/14.<sup>45</sup> Whether stabilization in glucose testing rates, as observed in this study, or preventive strategies and health initiatives have contributed to this decline requires further investigation. Regardless, our results suggest that further progress in implementing clinical practice guidelines could still be made.

Diabetes incidence has increased substantially among Chinese Canadian people, and a recent study found that immigrants in Ontario, especially South Asian and Southeast Asian people, convert from prediabetes to diabetes at earlier ages than people born in Canada.<sup>46,47</sup> Combined with reports of undiagnosed diabetes contributing about 20% to overall type 2 diabetes prevalence rates, continued vigilance in diabetes screening is needed.<sup>14</sup> In this study, we identified younger men (especially in the Chinese and general population groups), older Chinese and South Asian people, residents of low-income neighbourhoods, and those without a regular primary care physician or with hypertension or cardiovascular disease as potential targets to improve earlier diagnosis of diabetes.

Regular screening of vulnerable or at-risk populations is important for providing opportunities for early diagnosis and initiation of interventions, especially when the trajectory of the development of diabetes can be rapid, as has been observed in different ethnic groups.<sup>47</sup> Earlier diagnosis, particularly among those with additional risk factors, such as hypertension and hyperlipidemia, could lead to earlier glucose control, thus delaying the onset of diabetic complications, including cardiovascular disease. However, improving screening for diabetes requires at-risk individuals to seek testing, physicians to order testing, and public health agencies and policy-makers to increase awareness of the need for testing.



**Figure 3:** Annual glucose testing rates among high-risk populations in Ontario 2008–2017: (A) people with hypertension, (B) people with hyperlipidemia and (C) people with cardiovascular disease. “Glucose testing” is defined as at least 1 glycosylated hemoglobin, plasma or serum glucose or oral glucose tolerance test and excludes people who received a diabetes diagnosis before the study year or were admitted to hospital during the study year. “Cardiovascular disease” is defined as a history of hospital admission for myocardial infarction, stroke or heart failure, or previous percutaneous coronary intervention or coronary artery bypass graft surgery.

## Limitations

A strength of this study is the availability of multiple sources of routinely collected data, enabling us to examine trends in glucose testing among almost the entire adult population of Ontario. Nonetheless, our inability to delineate the reasons for testing is a limitation, as we recognize that some testing may be performed for reasons other than preventive care. Consequently, we used the term “glucose testing” rather than “diabetes screening” because we are unable to determine from our data sources whether a particular laboratory test was conducted for diabetes screening or another purpose, even though we considered only tests ordered for patients who were not in hospital and not pregnant.

Our analyses by ethnicity were also limited in that we could identify people who were only likely to be Chinese and South Asian, and although they are the 2 most populous ethnic groups (> 50% of all visible minorities) in Ontario, an investigation of other ethnic or racial groups was not possible.<sup>48</sup> In our multivariable analyses to identify characteristics associated with up-to-date testing, we were unable to account for some factors that may be associated with testing, such as education, obesity and physical activity. Finally, we examined primary care rostering post hoc and did not include it a priori in our analysis of predictors of up-to-date testing. However, we did include visit frequency and access to primary care physicians in these analyses.

## Conclusion

Rates of up-to-date glucose testing as recommended by Diabetes Canada guidelines remained relatively stable between 2010 and 2017, at about 70%. However, lower rates among some age, sex and ethnic groups and among people with hypertension or hyperlipidemia are concerning, given the morbidity and mortality associated with diabetes if left undiagnosed and untreated. Although much effort has been made in addressing rising diabetes prevalence and incidence, only with appropriate diabetes screening can indicated diabetes management strategies be implemented for those who receive a diagnosis or who are determined to be at risk. Improving screening rates will require initiatives to ensure that health practices and policies to encourage testing reach underscreened populations such that they are enabled to receive testing.

## References

1. Grundy SM, Benjamin IJ, Burke GL, et al. Diabetes and cardiovascular disease: a statement for healthcare professionals from the American Heart Association. *Circulation* 1999;100:1134-46.
2. 2011 Diabetes in Canada 2011: facts and figures from a public health perspective. Ottawa: Public Health Agency of Canada; 2011. Available: <https://www.phac-aspc.gc.ca/cd-mc/publications/diabetes-diabete/facts-figures-faits-chiffres-2011/pdf/facts-figures-faits-chiffres-eng.pdf> (accessed 2021 Jan. 13).
3. Global report on diabetes. Geneva: World Health Organization; 2016. Available: <https://www.who.int/publications/i/item/9789241565257> (accessed 2022 Feb. 14).
4. Geiss LS, Wang J, Cheng YJ, et al. Prevalence and incidence trends for diagnosed diabetes among adults aged 20 to 79 years, United States, 1980–2012. *JAMA* 2014;312:1218-26.
5. Houlden RL. Diabetes Canada 2018 Clinical Practice Guidelines for the Prevention and Management of Diabetes in Canada: Introduction. *Can J Diabetes* 2018; 42(Suppl 1):S1-5.
6. Selvin E, Parrinello CM, Sacks DB, et al. Trends in prevalence and control of diabetes in the United States, 1988–1994 and 1999–2010. *Ann Intern Med* 2014;160:517-25.
7. Boulé NG, Haddad E, Kenny GP, et al. Effects of exercise on glycemic control and body mass in type 2 diabetes mellitus: a meta-analysis of controlled clinical trials. *JAMA* 2001;286:1218-27.
8. Knowler WC, Barrett-Connor E, Fowler SE, et al.; Diabetes Prevention Program Research Group. Reduction of the incidence of type 2 diabetes with lifestyle intervention or metformin. *N Engl J Med* 2002;346:393-403.
9. Hoerger TJ, Hicks KA, Sorensen SW, et al. Cost-effectiveness of screening for pre-diabetes among overweight and obese US adults. *Diabetes Care* 2007; 30:2874-9.
10. Kahn R, Alperin P, Eddy D, et al. Age at initiation and frequency of screening to detect type 2 diabetes: a cost-effectiveness analysis. *Lancet* 2010;375:1365-74.
11. Wilson SE, Rosella LC, Lipscombe LL, et al. The effectiveness and efficiency of diabetes screening in Ontario, Canada: a population-based cohort study. *BMC Public Health* 2010;10:506.
12. Harris MI, Klein R, Welborn TA, et al. Onset of NIDDM occurs at least 4–7 yr before clinical diagnosis. *Diabetes Care* 1992;15:815-9.
13. Samuels TA, Cohen D, Brancati FL, et al. Delayed diagnosis of incident type 2 diabetes mellitus in the ARIC study. *Am J Manag Care* 2006;12:717-24.
14. Rosella LC, Lebenbaum M, Fitzpatrick T, et al. Prevalence of prediabetes and undiagnosed diabetes in Canada (2007–2011) according to fasting plasma glucose and HbA1c screening criteria. *Diabetes Care* 2015;38:1299-305.
15. Ekoé J-M, Goldenberg R, Katz P. Diabetes Canada 2018 Clinical Practice Guidelines for the Prevention and Management of Diabetes in Canada: Screening for diabetes in adults. *Can J Diabetes* 2018;42(Suppl 1):S16-9.
16. Tobe SW, Stone JA, Anderson T, et al. Canadian Cardiovascular Harmonized National Guidelines Endeavour (C-CHANGE) guideline for the prevention and management of cardiovascular disease in primary care: 2018 update. *CMAJ* 2018;190:E1192-206.
17. Creatore MI, Booth GL, Manuel DG, et al. A population-based study of diabetes incidence by ethnicity and age: support for the development of ethnic-specific prevention strategies. *Can J Diabetes* 2020;44:394-400.
18. Lipscombe LL, Hux JE. Trends in diabetes prevalence, incidence, and mortality in Ontario, Canada 1995–2005: a population-based study. *Lancet* 2007; 369:750-6.
19. Menke A, Casagrande S, Geiss L, et al. Prevalence of and trends in diabetes among adults in the United States, 1988–2012. *JAMA* 2015;314:1021-9.
20. Xu G, Liu B, Sun Y, et al. Prevalence of diagnosed type 1 and type 2 diabetes among US adults in 2016 and 2017: population based study. *BMJ* 2018;362:k1497.
21. Zghebi SS, Steinke DT, Carr MJ, et al. Examining trends in type 2 diabetes incidence, prevalence and mortality in the UK between 2004 and 2014. *Diabetes Obes Metab* 2017;19:1537-45.
22. Lipscombe LL, Hwee J, Webster L, et al. Identifying diabetes cases from administrative data: a population-based validation study. *BMC Health Serv Res* 2018;18:316.
23. Discharge Abstract Database metadata (DAD). Ottawa: Canadian Institute for Health Information. Available: <https://www.cihi.ca/en/discharge-abstract-database-metadata-dad> (accessed 2021 Jan. 11).
24. Schedule of benefits for laboratory services. Toronto: Ministry of Health, Ontario Health Insurance Plan, Laboratories and Genetics Branch; 2020. Available: <https://www.health.gov.on.ca/en/pro/programs/ohip/sob/> (accessed 2021 Jan. 11).
25. Tu K, Campbell NR, Chen Z-L, et al. Accuracy of administrative databases in identifying patients with hypertension. *Open Med* 2007;1:e18-26.
26. Tu JV, Chu A, Donovan LR, et al. The Cardiovascular Health in Ambulatory Care Research Team (CANHEART): using big data to measure and improve cardiovascular health and healthcare services. *Circ Cardiovasc Qual Outcomes* 2015;8:204-12.
27. Shah BR, Chiu M, Amin S, et al. Surname lists to identify South Asian and Chinese ethnicity from secondary data in Ontario, Canada: a validation study. *BMC Med Res Methodol* 2010;10:42.
28. Anderson K, Ross HJ, Austin PC, et al. Health care use before first heart failure hospitalization: identifying opportunities to pre-emptively diagnose impending decompensation. *JACC Heart Fail* 2020;8:1024-34.
29. Glazier RH, Klein-Geltink J, Kopp A, et al. Capitation and enhanced fee-for-service models for primary care reform: a population-based evaluation. *CMAJ* 2009;180:E72-81.
30. Online access to COVID-19 vaccination information and COVID-19 lab test results. Toronto: Ontario Laboratories Information System. Available: <https://www.ehealthontario.on.ca/en/for-healthcare-professionals/ontario-laboratories-information-system-olis> (accessed 2021 Jan. 11).
31. Donio PJ, Freitas C, Austin PC, et al. Comparison of readmission and death among patients with cardiac disease in Northern vs Southern Ontario. *Can J Cardiol* 2019;35:341-51.
32. Tu JV, Chu A, Rezai MR, et al. The incidence of major cardiovascular events in immigrants to Ontario, Canada: the CANHEART Immigrant Study. *Circulation* 2015;132:1549-59.
33. Canadian Diabetes Association Clinical Practice Guidelines Expert Committee; Ekoé J-M, Punthakee Z, Ransom T, et al. Screening for type 1 and type 2 diabetes. *Can J Diabetes* 2013;37(Suppl 1):S12-5.
34. Diabetes Canada Clinical Practice Guidelines Expert Committee; Meneilly GS, Knip A, Miller DB, et al. Diabetes in older people. *Can J Diabetes* 2018;42(Suppl 1):S283-95.



35. Creatore MI, Booth GL, Manuel DG, et al. Diabetes screening among immigrants: a population-based urban cohort study. *Diabetes Care* 2012;35:754-61.
36. Wilson SE, Lipscombe LL, Rosella LC, et al. Trends in laboratory testing for diabetes in Ontario, Canada 1995–2005: a population-based study. *BMC Health Serv Res* 2009;9:41.
37. Lee TM, Tobe SW, Butt DA, et al. Measuring cardiovascular quality in primary care using Canadian Cardiovascular Harmonization of National Guidelines Endeavour and electronic medical record data in Ontario. *CJCA Open* 2019;1:1-9.
38. Nichols J, Shah BR, Pequeno P, et al. Impact of a comprehensive guideline dissemination strategy on diabetes diagnostic test rates: an interrupted time series. *J Gen Intern Med* 2020;35:2662-7.
39. Ng E, Omariba DWR. Immigration, generational status and health literacy in Canada. *Health Educ J* 2014;73:668-82.
40. Tung EL, Baig AA, Huang ES, et al. Racial and ethnic disparities in diabetes screening between Asian Americans and other adults: BRFSS 2012–2014. *J Gen Intern Med* 2017;32:423-9.
41. Casagrande SS, Cowie CC, Genuth SM. Self-reported prevalence of diabetes screening in the U.S., 2005–2010. *Am J Prev Med* 2014;47:780-7.
42. Sheehy A, Pandhi N, Coursin DB, et al. Minority status and diabetes screening in an ambulatory population. *Diabetes Care* 2011;34:1289-94.
43. Magliano DJ, Islam RM, Barr ELM, et al. Trends in incidence of total or type 2 diabetes: systematic review. *BMJ* 2019;366:15003.
44. Magliano DJ, Chen L, Islam RM, et al. Trends in the incidence of diagnosed diabetes: a multicountry analysis of aggregate data from 22 million diagnoses in high-income and middle-income settings. *Lancet Diabetes Endocrinol* 2021;9:203-11.
45. Canadian Chronic Disease Surveillance System (CCDSS), Data Tool 2000–2017, 2019 Edition. Ottawa: Public Health Agency of Canada; modified 2021 Dec. 15. Available: <https://health-infobase.canada.ca/ccdss/Index> (accessed 2021 Jan. 11).
46. Alangh A, Chiu M, Shah BR. Rapid increase in diabetes incidence among Chinese Canadians between 1996 and 2005. *Diabetes Care* 2013;36:3015-7.
47. Fazli GS, Moineddin R, Bierman AS, et al. Ethnic variation in the conversion of prediabetes to diabetes among immigrant populations relative to Canadian-born residents: a population-based cohort study. *BMJ Open Diabetes Res Care* 2020;8:e000907.
48. Data tables, 2016 Census. Ottawa: Statistics Canada; modified 2019 June 17. Available: <https://www12.statcan.gc.ca/census-recensement/2016/dp-pd/dt-td/Rp-eng.cfm?TABID=2&Lang=E&APATH=3&DETAIL=0&DIM=0&FL=A&FREE=0&GC=0&GID=1341679&GK=0&GRP=1&PID=110531&PRID=10&PTYPE=109445&S=0&SHOWALL=0&SUB=0&Temporal=2017&TH EME=120&VID=0&VNAMEE=&VNAM EF=&D1=0> (accessed 2021 Jan. 11).

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**Data sharing:** The data set from this study is held securely in coded form at ICES. While data sharing agreements prohibit ICES from making the data set publicly available, access may be granted to those who meet prespecified criteria for confidential access, available at <https://www.ices.on.ca/DAS>.

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