

# Research

# Depression, diabetes and immigration status: a retrospective cohort study using the Canadian Longitudinal Study on Aging

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# **Abstract**

**Background:** A bidirectional association between depression and diabetes exists, but has not been evaluated in the context of immigrant status. Given that social determinants of health differ between immigrants and nonimmigrants, we evaluated the association between diabetes and depression incidence, depression and diabetes incidence, and whether immigrant status modified this association, among immigrants and nonimmigrants in Canada.

**Methods:** We employed a retrospective cohort design using data from the Canadian Longitudinal Study on Aging Comprehensive cohort (baseline [2012–2015] and 3-year follow-up [2015–2018]). We defined participants as having diabetes if they self-reported it or if their glycated hemoglobin A<sub>1c</sub> level was 7% or more; we defined participants as having depression if their Center for Epidemiological Studies Depression score was 10 or higher or if they were currently undergoing depression treatment. We excluded those with baseline depression (Cohort 1) and baseline diabetes (Cohort 2) to evaluate the associations between diabetes and depression incidence, and between depression and diabetes incidence, respectively. We constructed logistic regression models with interaction by immigrant status.

**Results:** Cohort 1 (n = 20.723; mean age 62.7 yr, standard deviation [SD] 10.1 yr; 47.6% female) included 3766 (18.2%) immigrants. Among immigrants, 16.4% had diabetes, compared with 15.6% among nonimmigrants. Diabetes was associated with an increased risk of depression in nonimmigrants (adjusted odds ratio [OR] 1.27, 95% confidence interval [CI] 1.08–1.49), but not in immigrants (adjusted OR 1.12, 95% CI 0.80–1.56). Younger age, female sex, weight change, poor sleep quality and pain increased depression risk. Cohort 2 (n = 22.054; mean age 62.1 yr, SD 10.1 yr; 52.2% female) included 3913 (17.7%) immigrants. Depression was associated with an increased risk of diabetes in both nonimmigrants (adjusted OR 1.39, 95% CI 1.16–1.68) and immigrants (adjusted OR 1.60, 95% CI 1.08–2.37). Younger age, male sex, waist circumference, weight change, hypertension and heart disease increased diabetes risk.

**Interpretation:** We found an overall bidirectional association between diabetes and depression that was not significantly modified by immigrant status. Screening for diabetes for people with depression and screening for depression for those with diabetes should be considered.

creening for diabetes and depression is recommended in recently arrived immigrants.<sup>1</sup> In addition, given some evidence for a bidirectional association between diabetes and depression in the general population,<sup>2-5</sup> ongoing surveillance for diabetes may be relevant in the context of depression and, likewise, ongoing surveillance for depression may be justified with people with diabetes, particularly among immigrants.

Immigrants may be particularly vulnerable to the development of diabetes and depression<sup>6-9</sup> owing to factors associated with resettlement, such as acculturation, stress and social and economic challenges.<sup>5,10–17</sup> For example, studies have reported that immigrants from South Asian countries are generally healthy upon arrival but rapidly develop diabetes after immigration.<sup>18,19</sup> Studies have also reported that

immigrants with diabetes are at higher risk of depression than the general host population<sup>6,7</sup> and the population in their country of origin. <sup>12,20–25</sup> Conversely, immigrants with depression may also be at high risk of diabetes because of complex social determinants of health <sup>1,18,19,26</sup> and delayed diagnosis of and treatment for diabetes; <sup>27–31</sup> use of antidepressants may also increase diabetes risk. <sup>32,33</sup>

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To explore the bidirectional association between diabetes and depression in the context of immigrant status, we evaluated the association between diabetes and depression incidence, the association between depression and diabetes incidence, and whether immigration status modified this association, among immigrants and nonimmigrants in Canada.

## **Methods**

#### Study design

We used a retrospective cohort design to evaluate the association between diabetes and depression incidence (Cohort 1) and the association between depression and diabetes incidence (Cohort 2).

We used data from the Canadian Longitudinal Study on Aging (CLSA) Comprehensive cohort at baseline (2012–2015) and 3-year follow-up (2015–2018) to construct our study cohorts.<sup>34,35</sup> The paper was reported according to the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) checklist.<sup>36</sup>

#### Data source and population

The CLSA Comprehensive cohort included community-dwelling individuals, aged 45–85 years, who were randomly selected from within a 25-km radius of 11 sites in cities across Canada using the provincial health care registration databases and random digit dialing of landline telephones.<sup>34</sup> Participants provided demographic, social, physical, clinical, psychological and economic data at baseline and at 3-year follow-up through an in-person interview, on-site physical examinations and blood and urine collection. Details about the CLSA's sampling, design and data collection have been published.<sup>34,35</sup>

We defined participants as having diabetes if they selfreported a diagnosis with any type of diabetes (i.e., answered yes to "Has a doctor ever told you that you have diabetes, borderline diabetes or that your blood sugar is high?") or if their glycated hemoglobin (HbA<sub>1c</sub>) level was 7% or more. We defined participants as having depression if their Center for Epidemiological Studies Depression-10 Scale (CES-D-10) score was 10 or higher or if they self-reported being currently treated for depression.<sup>37–40</sup> For both Cohort 1 and Cohort 2, we excluded participants missing information on immigration, depression or diabetes at baseline. In addition, for Cohort 1, we excluded those with baseline depression and those with missing depression status in follow-up; for Cohort 2, we excluded those with baseline diabetes and those with missing diabetes status in follow-up. The CLSA defined immigrants using 2 questions: "In what country were you born?" and "In what year did you first come to Canada to live?"41

#### **Exposure and outcomes**

Our outcomes were depression incidence for Cohort 1 and diabetes incidence for Cohort 2, assessed using the 3-year follow-up data. Depression was defined the same at follow-up as at baseline, but diabetes in follow-up was based only on self-report because  $HbA_{1c}$  data were unavailable at that time point. The main exposure variables were baseline diabetes for Cohort 1 and baseline depression for Cohort 2.

#### **Baseline characteristics**

The baseline characteristics for Cohort 1 and Cohort 2 included those previously associated with both diabetes and depression, namely self-reported predisposing sociodemographic and socioeconomic factors, medical conditions and lifestyle choices, as well as anthropomorphic measurements and blood assessments. The predisposing sociodemographic and socioeconomic factors included sex, age, ethnic or racial background, immigration status, time since migration, marital status, language most spoken at home, household income, employment status, education, province and place of residence. The place of residence classification (urban or rural) was derived by the CLSA based on Statistics Canada's Postal Code Conversion File, whereby rural areas were those with a total population of fewer than 10 000 people.<sup>42</sup> The medical conditions included pain, 43 cancer, 44 arthritis, 45 bowel disorders 46 and hypertension.<sup>47</sup> The lifestyle choices included perceived health, sleep satisfaction, alcohol consumption, smoking status and nutritional risk. The anthropomorphic measurements and blood assessments included body mass index, waist circumference,<sup>48</sup> weight change<sup>49</sup> and vitamin D deficiency.<sup>50,51</sup> Details on these baseline characteristics are provided in Appendix 1, available at www.cmajopen.ca/content/10/2/E508/suppl/DC1.

In addition, we assessed self-reported comorbidities associated with depression, including heart disease,<sup>52,53</sup> kidney disease<sup>54</sup> and iron deficiency,<sup>50,51,55</sup> for Cohort 1; we considered participant lipid profiles (assessed from blood tests)<sup>56</sup> and lean and fat mass (measured with dual-energy x-ray absorptiometry)<sup>57</sup> for Cohort 2.

### Statistical analysis

For each cohort, we computed descriptive statistics with means and standard deviations (SDs) for continuous variables and counts with percentages for categorical variables, by immigration and baseline diabetes statuses (Cohort 1) or by immigration and baseline depression statuses (Cohort 2).

Among immigrants and nonimmigrants, we compared baseline characteristics by baseline diabetes status (Cohort 1) or by baseline depression status (Cohort 2) using multivariable logistic regression models. We used these models to evaluate the associations between baseline diabetes and the risk of depression at 3 years in Cohort 1 and between depression at baseline and the risk of 3-year diabetes in Cohort 2. To assess the effect modification by immigration status, we included interaction terms for diabetes and immigration status and for depression and immigration status in the models for Cohort 1 and Cohort 2, respectively.

The baseline characteristics listed above were considered for inclusion in the multivariable model. Immigration status, sex and age were forced in all models. We removed other variables that were not significant (p > 0.5), did not affect the Bayesian Information Criterion upon inclusion and did not modify the effect of the main exposure variable by more than 10%. To make the estimates generalizable to the Canadian population, we used the CLSA analytical sample weights and geographic strata information in the regression analyses.  $^{60,61}$  Results were expressed in odds ratios (ORs) and 95% confidence intervals (CIs).





We also conducted sensitivity analyses. First, we defined baseline diabetes solely by the self-reported physician diagnosis and repeated the main analyses for Cohort 1. Second, we conducted the main analyses excluding individuals with missing variables. We repeated these analyses using multiple imputations to impute missing information using the Markov Chain Monte Carlo method, as implemented in SAS Proc MI.<sup>62</sup> Five imputed data sets were generated based on all baseline characteristics. The results from these data sets were combined using Rubin's rules,<sup>63</sup> as implemented by SAS Proc MIANALYZE. We performed all statistical analyses using SAS version 9.4 (SAS Institute).

#### Ethics approval

Ethics approval was provided by the McGill University Health Centre Research Ethics Board.

#### Results

#### Diabetes at baseline and risk of depression at 3 years

Cohort 1 included 20723 individuals (Figure 1), including 3766 (18.2%) immigrants. Among immigrants, 616 (16.4%) had diabetes at baseline, compared with 2639 (15.6%) among

nonimmigrants (Table 1). In general, baseline characteristics that differed between individuals with or without baseline diabetes were similar in immigrants and nonimmigrants (Appendix 1, Table 1).

Overall, 339 (10.4%) of people with diabetes had depression at 3 years, compared with 1465 (8.4%) people without diabetes. Among immigrants and nonimmigrants, respectively, 60 (9.7%) and 279 (10.6%) of those with baseline diabetes had depression at 3 years, compared with 265 (8.4%) and 1200 (8.4%), respectively, of those without baseline diabetes (Table 2).

In multivariable logistic regression models (Table 2), individuals with baseline diabetes had 18% higher odds of depression at 3 years than those without baseline diabetes (adjusted OR 1.18, 95% CI 1.01–1.37). Among nonimmigrants, baseline diabetes was associated with 27% increased odds of depression at 3 years (adjusted OR 1.27, 95% CI 1.08–1.49), whereas no significant association was observed among immigrants (adjusted OR 1.12, 95% CI 0.80–1.56).

Depression at 3 years was associated with being female, living with pain, current smoking (v. past or never), weight change in the previous year (loss or gain), living in Quebec (v. Ontario or British Columbia), being younger (45–60 v. 61–70 yr) and not being satisfied with sleep (Appendix 1, Table 3).

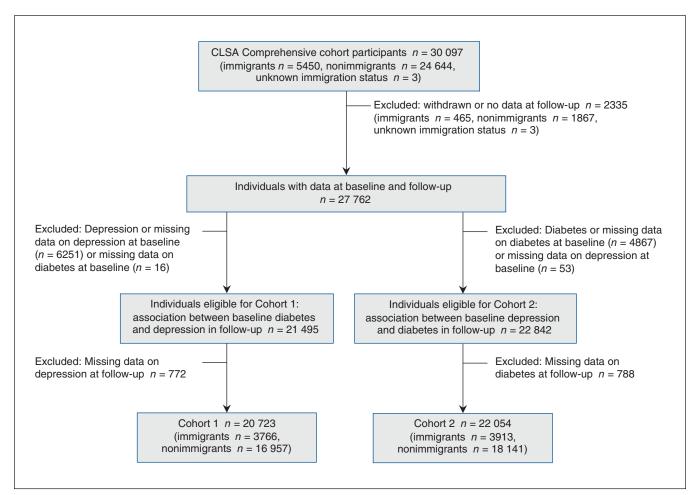


Figure 1: Flow chart of study cohorts. Note: CLSA = Canadian Longitudinal Study on Aging.



	Immi	igrant	Nonimmigrant		
Characteristic	No. (%) of participants with diabetes $n = 616$	No. (%) of participants without diabetes $n = 3150$	No. (%) of participants with diabetes $n = 2639$	No. (%) of participants without diabetes n = 14318	
Age, yr					
45–60	141 (22.9)	1173 (37.2)	893 (33.8)	7050 (49.2)	
61–70	236 (38.3)	1050 (33.3)	969 (36.7)	4224 (29.5)	
71–85	239 (38.8)	927 (29.4)	777 (29.4)	3044 (21.3)	
Sex	200 (00.0)	<i>021</i> (20.4)	777 (20.4)	0044 (21.0)	
Male	408 (66.2)	1743 (55.3)	1551 (58.8)	7167 (50.1)	
Female	208 (33.8)	1407 (44.7)	1088 (41.2)	7151 (49.9)	
Marital status	200 (00.0)	1407 (44.7)	1000 (41.2)	7131 (49.9)	
Single	27 (4.4)	149 (4.7)	234 (8.9)	1176 (8.2)	
Married	455 (73.9)	. ,	1834 (69.5)	. ,	
Widowed, divorced or separated	134 (21.8)	2405 (76.3)	571 (21.6)	10 407 (72.7)	
Missing	104 (∠ 1.8)	595 (18.9)	3/1 (21.0)	2731 (19.1)	
Language most spoken at home	<del>-</del>	1 (0.0)	<del>-</del>	4 (0.0)	
	21 /5 0)	00F (C F)	E41 (00 E)	2171 (00.1)	
French	31 (5.0)	205 (6.5)	541 (20.5)	3171 (22.1)	
English	492 (79.9)	2597 (82.4)	2092 (79.3)	11 114 (77.6)	
Other	55 (8.9)	254 (8.1)	3 (0.1)	17 (0.2)	
Missing	38 (6.2)	94 (3.0)	3 (0.1)	16 (0.1)	
Ethnic or racial background	404 (75.0)	0040 (00.0)	0500 (00.4)	11100 (00.0)	
White	464 (75.3)	2640 (83.8)	2596 (98.4)	14 166 (98.9)	
Black	36 (5.8)	89 (2.8)	8 (0.3)	19 (0.1)	
South Asian	50 (8.1)	122 (3.9)	0 (0.0)	6 (0.0)	
Chinese	24 (3.9)	107 (3.4)	8 (0.3)	37 (0.3)	
Other	42 (6.8)	188 (6.0)	25 (0.9)	80 (0.6)	
Missing	_	4 (0.1)	2 (0.1)	10 (0.1)	
Years since arrival to Canada					
< 20	42 (6.8)	396 (12.6)	_	_	
20–40	162 (26.3)	781 (24.8)	<del>-</del> -	_	
> 40	412 (66.9)	1973 (62.6)	_	_	
Total household income, \$					
< 20 000	29 (4.7)	82 (2.6)	140 (5.3)	432 (3.0)	
20 000-50 000	141 (22.9)	589 (18.7)	641 (24.3)	2435 (17.0)	
50 000-100 000	245 (39.8)	1076 (34.2)	927 (35.1)	4786 (33.4)	
> 100 000	163 (26.5)	1180 (37.5)	779 (29.5)	5855 (40.9)	
Missing	38 (6.1)	223 (7.0)	152 (5.8)	810 (5.7)	
Working status					
Employed	158 (25.6)	1333 (42.3)	826 (31.3)	6368 (44.5)	
Unemployed	29 (4.7)	111 (3.5)	72 (2.7)	468 (3.3)	
Retired	429 (69.6)	1699 (54.0)	1732 (65.6)	7453 (52.1)	
Missing	_	7 (0.2)	9 (0.3)	29 (0.2)	
Education level					
Less than secondary school	20 (3.2)	77 (2.4)	188 (7.1)	590 (4.1)	
Secondary school	51 (8.3)	195 (6.2)	316 (12.0)	1275 (8.9)	
Postsecondary degree or diploma	541 (87.8)	2870 (91.1)	2134 (80.9)	12436 (86.9)	
Missing	4 (0.6)	8 (0.3)	1 (0.0)	17 (0.1)	



	Immi	igrant	Nonimmigrant		
Characteristic	No. (%) of participants with diabetes $n = 616$	No. (%) of participants without diabetes $n = 3150$	No. (%) of participants with diabetes $n = 2639$	No. (%) of participants without diabetes $n = 14318$	
Place of residence*					
Rural	58 (9.4)	310 (9.8)	365 (13.8)	1849 (12.9)	
Urban	551 (89.4)	2799 (88.9)	2239 (84.8)	12303 (85.9)	
Missing	7 (1.1)	41 (1.3)	35 (1.4)	166 (1.2)	
Province	, ()	11 ()	30 (,	100 ()	
Quebec	65 (10.6)	365 (11.6)	536 (20.3)	3199 (22.3)	
British Columbia	189 (30.7)	996 (31.6)	538 (20.4)	2736 (19.1)	
Ontario	169 (27.4)	869 (27.6)	551 (20.9)	2920 (20.4)	
Other†	193 (31.3)	920 (29.2)	1014 (38.4)	5463 (38.2)	
Medical conditions‡	190 (01.0)	320 (20.2)	1014 (00.7)	J400 (50.2)	
Living with pain	226 (36.7)	944 (30.0)	1055 (40.0)	4260 (29.8)	
Bowel disorders	41 (6.7)	227 (7.2)	224 (8.5)	1135 (7.9)	
Arthritis	. ,	83 (2.6)	109 (4.1)		
Heart disease	16 (2.6)	1007 (32.0)	1552 (58.8)	355 (2.5)	
	370 (60.1)		. ,	4353 (30.4)	
Kidney disease	36 (5.8)	66 (2.1)	111 (4.2)	292 (2.0)	
Stroke	11 (1.8)	36 (1.1)	65 (2.5)	157 (1.1)	
Cancer	95 (15.4)	475 (15.1)	465 (17.6)	2005 (14.0)	
Hypertension	355 (57.6)	1081 (34.3)	1530 (58.0)	4851 (33.9)	
Anxiety disorder	17 (2.8)	109 (3.5)	137 (5.2)	606 (4.2)	
Alcohol consumption	00 (14.0)	000 (0.0)	0.40 (40.0)	1000 (0.4)	
Never	92 (14.9)	309 (9.8)	340 (12.9)	1209 (8.4)	
About once a month	141 (22.9)	514 (16.3)	608 (23.0)	2255 (15.7)	
2–4 times a month	115 (18.7)	576 (18.3)	588 (22.3)	3160 (22.1)	
> 2 times a week	239 (38.8)	1641 (52.1)	1031 (39.1)	7433 (51.9)	
Missing	29 (4.7)	110 (3.5)	72 (2.7)	261 (1.8)	
Smoking status	(40.0)	(50.5)	(44.0)	= := t (=0.0)	
Nonsmoker	297 (48.2)	1691 (53.7)	1088 (41.2)	7184 (50.2)	
Former smoker	275 (44.6)	1291 (41.0)	1348 (51.1)	6029 (42.1)	
Smoker	44 (7.1)	168 (5.3)	203 (7.7)	1105 (7.7)	
Sleep quality		45.5.1			
Satisfied or very satisfied	373 (60.6)	2014 (63.9)	1615 (61.2)	9114 (63.7)	
Neutral	116 (18.8)	513 (16.3)	388 (14.7)	2158 (15.1)	
Dissatisfied or very dissatisfied	127 (20.6)	617 (19.6)	635 (24.1)	3038 (21.2)	
Missing		6 (0.2)	1 (0.0)	8 (0.1)	
Nutritional risk status					
Low risk	394 (64.0)	2256 (71.6)	1577 (59.8)	10 128 (70.7)	
High risk	201 (32.6)	796 (25.3)	1008 (38.2)	3919 (27.4)	
Missing	21 (3.4)	98 (3.1)	54 (2.0)	271 (1.9)	
Weight classification§					
Normal weight	112 (18.2)	1193 (37.9)	383 (14.5)	4827 (33.7)	
Overweight	248 (40.3)	1312 (41.7)	967 (36.6)	6086 (42.5)	
Obese	253 (41.1)	638 (20.3)	1277 (48.4)	3365 (23.5)	
Missing	3 (0.5)	7 (0.2)	12 (0.5)	40 (0.3)	

<sup>\*</sup>The place of residence classification (urban or rural) was derived by CLSA based on Statistics Canada's Postal Code Conversion File, which defines rural areas as those with a total population of fewer than 10 000 people.

<sup>†</sup>Other provinces are Alberta, Manitoba, Nova Scotia, Prince Edward Island, Newfoundland and Labrador, and Saskatchewan.

<sup>#</sup>Missing data on medical conditions: living with pain (n = 262, 1.3%), bowel disorders (n = 49, 0.2%), arthritis (n = 168, 0.8%), heart disease (n = 27, 0.1%), kidney disease (n = 41, 0.2%), stroke (n = 35, 0.2%), cancer (n = 19, 0.1%), hypertension (n = 2472, 11.9%), anxiety disorder (n = 25, 0.1%). §Based on body mass index international classification for adults aged  $\geq 18$  years.



	No. (%) of participants			Weighted OR (95% CI)*		
•		No				
Variable	Depression	depression	Total	Crude	Adjusted	
Immigrant						
Diabetes	60 (9.7)	556 (90.3)	616	1.19 (0.86–1.65)		
No diabetes	265 (8.4)	2885 (91.6)	3150			
Nonimmigrant						
Diabetes	279 (10.6)	2360 (89.4)	2639	1.35 (1.15–1.57)		
No diabetes	1200 (8.4)	13 035 (91.6)	14318			
Diabetes v. no diabetes					1.18 (1.01–1.37)	
Interaction effect of immigration status and diabetes at baseline						
Immigrant with diabetes v. immigrant without diabetes					1.12 (0.80–1.56	
Immigrant with diabetes v. nonimmigrant with diabetes					1.01 (0.72–1.41	
Immigrant without diabetes v. nonimmigrant without diabetes					1.15 (0.98–1.35	
Nonimmigrant with diabetes v. nonimmigrant without diabetes					1.27 (1.08–1.49	

Note: CI = confidence interval, OR = odds ratio.

\*The model included all the variables listed in Appendix 1, Table 1. Immigration status, sex and age were forced in all models. We removed other variables that were not significant ( $\rho > 0.5$ ), which did not affect the Bayesian Information Criterion upon inclusion and which did not modify the effect of the main exposure variable by more than 10%. Survey weights were provided by the Canadian Longitudinal Study on Aging. The full model is available in Appendix 1, Table 2.

## Depression at baseline and risk of diabetes at 3 years

Cohort 2 included 22 054 individuals (Figure 1), including 3913 (17.7%) immigrants (Table 3). Among immigrants, 762 (19.5%) had depression at baseline, compared with 3797 (20.9%) non-immigrants. In general, baseline characteristics between individuals with or without baseline depression were similar in immigrants and in nonimmigrants (Appendix 1, Table 4).

Overall, 311 (6.8%) individuals with depression at baseline developed diabetes, compared with 837 (4.8%) of those without depression (Table 4). Among immigrants and nonimmigrants, respectively, 54 (7.1%) and 257 (6.8%) of those with depression at baseline developed diabetes, compared with 144 (4.6%) and 693 (4.8%) of those without depression at baseline.

In multivariable logistic regression models (Table 4), those with depression at baseline had 43% higher odds to develop diabetes than those without depression (adjusted OR 1.43, 95% CI 1.21–1.68). Among nonimmigrants, depression was associated with 39% increased odds of diabetes (adjusted OR 1.39, 95% CI 1.16–1.68); among immigrants, depression was associated with a 60% increased odds of diabetes (adjusted OR 1.60, 95% CI 1.08–2.37).

Overall, diabetes at 3 years was associated with being male, being younger (45–60 v. 71–85 yr), having hypertension, having heart disease, high waist circumference and weight change (v. same weight) (Appendix 1, Table 3).

Results of both sensitivity analyses were similar to those of the main analyses (Appendix 1, Table 2, Table 5, Table 6).

## Interpretation

Our study provides evidence for a bidirectional association between diabetes and depression in the population aged 45 years and older. Overall, diabetes at baseline was associated with an increased risk of depression at 3-year followup, and depression at baseline was associated with an increased risk of diabetes at 3-year follow-up. In general, immigration status did not modify these risks in either direction. Specifically, although depression was associated with 39% and 60% increased odds of diabetes among nonimmigrants and immigrants, respectively, the overlapping 95% CIs did not suggest a conclusive modifying effect by immigration status. Moreover, although diabetes at baseline was associated with 27% increased odds of depression among nonimmigrants and a nonsignificant increase by 12% among immigrants, a modification effect by immigration status could not be concluded.

We had expected to observe differences in the bidirectional relation between diabetes and depression by immigration status because of the complex social determinants of health and stressors<sup>29</sup> that immigrants live with and how these may potentially be associated with low-grade inflammation.<sup>64-66</sup> The lack of effect modification by immigration status in the diabetes–depression relation in our study may perhaps be explained by the resilience of immigrants in their dynamic process of positive adaptation.<sup>24,67</sup>



	Immi	grant	Nonimmigrant		
Characteristic	No. (%) of participants with depression $n = 762$	No. (%) of participants without depression $n = 3151$	No. (%) of participants with depression $n = 3797$	No. (%) of participants without depression $n = 14344$	
Age, yr					
45–60	325 (42.7)	1173 (37.2)	1941 (51.1)	7053 (49.2)	
61–70	235 (30.8)	1048 (33.3)	1087 (28.6)	4228 (29.5)	
71–85	202 (26.5)	930 (29.5)	769 (20.3)	3063 (21.4)	
Sex	()				
Male	266 (34.9)	1741 (55.3)	1351 (35.6)	7175 (50.0)	
Female	496 (65.1)	1410 (44.7)	2446 (64.4)	7169 (50.0)	
Marital status	400 (00.1)	1410 (44.7)	2440 (04.4)	7 100 (00.0)	
Single	68 (8.9)	149 (4.7)	485 (12.8)	1178 (8.2)	
Married	465 (61.0)	2403 (76.3)	2223 (58.5)	10417 (72.6)	
Widowed, divorced or separated	227 (29.8)	598 (19.0)	1088 (28.7)	2745 (19.1)	
Missing	· · · · · · · · · · · · · · · · · · ·	1 (0.0)	1 (0.0)	4 (0.0)	
Language most spoken at home	2 (0.3)	1 (0.0)	1 (0.0)	4 (0.0)	
	GE (9.E)	205 (6.5)	005 (04.4)	2174 (22.1)	
French	65 (8.5)	205 (6.5)	925 (24.4)	3174 (22.1)	
English	597 (78.3)	2598 (82.5)	2861 (75.3)	11 137 (77.6)	
Other	58 (7.6)	254 (8.1)	7 (0.2)	17 (0.1)	
Missing	42 (5.5)	94 (3.0)	4 (0.1)	16 (0.1)	
Ethnic or racial background	200 (00 =)	2244 (22.2)			
White	629 (82.5)	2641 (83.8)	3758 (99.0)	14 191 (98.9)	
Black	24 (3.1)	89 (2.8)	8 (0.2)	19 (0.1)	
South Asian	36 (4.7)	122 (3.9)	2 (0.1)	6 (0.0)	
Chinese	17 (2.2)	107 (3.4)	7 (0.2)	37 (0.3)	
Other	53 (7.0)	188 (6.0)	20 (0.5)	81 (0.6)	
Missing	3 (0.4)	4 (0.1)	2 (0.1)	10 (0.1)	
Years since arrival to Canada					
< 20	112 (14.7)	395 (12.5)	_		
20–40	190 (24.9)	783 (24.8)	_	_	
> 40	460 (60.4)	1973 (62.6)	<u>-</u>		
Total household income, \$					
< 20 000	57 (7.5)	82 (2.6)	332 (8.7)	436 (3.0)	
20 000–50 000	208 (27.3)	591 (18.8)	923 (24.3)	2443 (17.0)	
50 000-100 000	246 (32.3)	1075 (34.1)	1217 (32.1)	4794 (33.4)	
> 100 000	194 (25.5)	1180 (37.4)	1088 (28.7)	5854 (40.8)	
Missing	57 (7.5)	223 (7.1)	237 (6.2)	817 (5.7)	
Working status					
Employed	297 (39.0)	1332 (42.3)	1510 (39.8)	6373 (44.4)	
Unemployed	71 (9.3)	111 (3.5)	313 (8.2)	468 (3.3)	
Retired	393 (51.6)	1701 (54.0)	1964 (51.7)	7474 (52.1)	
Missing	1 (0.1)	7 (0.2)	10 (0.3)	29 (0.2)	
Education level					
Less than secondary school	30 (3.9)	76 (2.4)	260 (6.8)	595 (4.1)	
Secondary school	64 (8.4)	196 (6.2)	367 (9.7)	1278 (8.9)	
Postsecondary degree or diploma	664 (87.1)	2871 (91.1)	3166 (83.4)	12454 (86.8)	
Missing	4 (0.5)	8 (0.3)	4 (0.1)	17 (0.1)	



	Imm	igrant	Nonimmigrant		
Characteristic	No. (%) of participants with depression $n = 762$	No. (%) of participants without depression $n = 3151$	No. (%) of participants with depression $n = 3797$	No. (%) of participants without depression $n = 14344$	
Place of residence*					
Rural	75 (9.8)	310 (9.8)	453 (11.9)	1850 (12.9)	
Urban	674 (88.5)	2800 (88.9)	3293 (86.7)	12328 (85.9)	
Missing	13 (1.7)	41 (1.3)	51 (1.3)	166 (1.2)	
Province	, ,				
Quebec	111 (14.6)	365 (11.6)	922 (24.3)	3204 (22.3)	
British Columbia	227 (29.8)	995 (31.6)	665 (17.5)	2735 (19.1)	
Ontario	223 (29.3)	872 (27.7)	855 (22.5)	2921 (20.4)	
Other†	201 (26.4)	919 (29.2)	1355 (35.7)	5484 (38.2)	
Medical conditions‡	V - /	( - )	\ /	\ /	
Living with pain	366 (48.0)	945 (30.0)	1904 (50.1)	4267 (29.7)	
Bowel disorders	105 (13.8)	227 (7.2)	582 (15.3)	1136 (7.9)	
Arthritis	28 (3.7)	83 (2.6)	160 (4.2)	354 (2.5)	
Heart disease	280 (36.7)	1009 (32.0)	1442 (38.0)	4364 (30.4)	
Kidney disease	17 (2.2)	67 (2.1)	109 (2.9)	294 (2.0)	
Stroke	15 (2.0)	37 (1.2)	67 (1.8)	157 (1.1)	
Cancer	117 (15.4)	474 (15.0)	573 (15.1)	2012 (14.0)	
Hypertension	283 (37.1)	1080 (34.3)	1457 (38.4)	4870 (34.0)	
Anxiety disorder	119 (15.6)	109 (3.5)	919 (24.2)	611 (4.3)	
Alcohol consumption	()		0.0 (2.02)		
Never	101 (13.3)	310 (9.8)	528 (13.9)	1218 (8.5)	
About once a month	171 (22.4)	512 (16.2)	756 (19.9)	2259 (15.7)	
2–4 times a month	150 (19.7)	574 (18.2)	827 (21.8)	3166 (22.1)	
> 2 times a week	310 (40.7)	1645 (52.2)	1625 (42.8)	7436 (51.8)	
Missing	30 (3.9)	110 (3.5)	61 (1.6)	265 (1.8)	
Smoking status	00 (0.0)	110 (0.0)	01 (1.0)	200 (1.0)	
Nonsmoker	377 (49.5)	1690 (53.6)	1644 (43.3)	7200 (50.2)	
Former smoker	315 (41.3)	1293 (41.0)	1690 (44.5)	6040 (42.1)	
Smoker	70 (9.2)	168 (5.3)	463 (12.2)	1104 (7.7)	
Sleep quality	. 0 (0.2)	100 (0.0)	100 (12.2)	1104 (1.1)	
Satisfied or very satisfied	339 (44.5)	2014 (63.9)	1692 (44.6)	9133 (63.7)	
Neutral	132 (17.3)	513 (16.3)	588 (15.5)	2162 (15.1)	
Dissatisfied or very dissatisfied	291 (38.2)	618 (19.6)	1512 (39.8)	3041 (21.2)	
Missing		6 (0.2)	5 (0.1)	8 (0.1)	
Nutritional risk status		U.L)	J (0.1)	J (0.1)	
Low risk	381 (50.0)	2256 (71.6)	1772 (46.7)	10 147 (70.7)	
High risk	356 (46.7)	797 (25.3)	1941 (51.1)	3924 (27.4)	
Missing		· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·		
Weight classification§	25 (3.3)	98 (3.1)	84 (2.2)	273 (1.9)	
Normal weight	264 (34.6)	110/ (270)	1205 (217)	/838 (33.7)	
Overweight	. ,	1194 (37.9)	1205 (31.7)	4838 (33.7)	
	310 (40.7)	1313 (41.7)	1423 (37.5)	6094 (42.5)	
Obese Missing	184 (24.1) 4 (0.5)	637 (20.2) 7 (0.2)	1144 (30.1) 25 (0.7)	3372 (23.5) 40 (0.3)	

<sup>\*</sup>The place of residence classification (urban or rural) was derived by CLSA based on Statistics Canada's Postal Code Conversion File, which defines rural areas as those

with a total population of fewer than 10 000 people. †Other provinces are Alberta, Manitoba, Nova Scotia, Prince Edward Island, Newfoundland and Labrador, and Saskatchewan. ‡Missing data on medical conditions: living with pain (n = 293, 1.3%), bowel disorders (n = 51, 0.2%), arthritis (n = 197, 0.9%), heart disease (n = 33, 0.1%), kidney disease (n = 44, 0.2%), stroke (n = 37, 0.2%), cancer (n = 25, 0.1%), hypertension (n = 2438, 11.0%), anxiety disorder (n = 44, 0.2%). §Based on body mass index international classification for adults aged  $\geq 18$  years.



	No. (%) of participants			Weighted OR (95% CI)*		
Variable	Diabetes	No diabetes	Total	Crude	Adjusted	
Immigrant						
Depression	54 (7.1)	708 (92.9)	762	1.66 (1.16–2.39)		
No depression	144 (4.6)	2970 (95.4)	3151			
Nonimmigrant						
Depression	257 (6.8)	3540 (93.2)	3797	1.55 (1.31–1.82)		
No depression	693 (4.8)	13 651 (95.2)	14 344			
Depression v. no depression					1.43 (1.21–1.68	
Interaction effect of immigrant status and depression at baseline						
Immigrant with depression v. immigrant without depression					1.60 (1.08–2.3	
Immigrant with depression v. nonimmigrant without depression					1.10 (0.76–1.58	
Immigrant without depression v. nonimmigrant without depression					0.96 (0.76–1.2	
Nonimmigrant with depression v. nonimmigrant without depression					1.39 (1.16–1.6	

Note: CI = confidence interval, OR = odds ratio.

\*The model included all the variables listed in Appendix 1, Table 3. Immigration status, sex and age were forced in all models. We removed other variables that were not significant ( $\rho > 0.5$ ), which did not affect the Bayesian Information Criterion upon inclusion and which did not modify the effect of the main exposure variable by more than 10%. Survey weights were provided by the Canadian Longitudinal Study on Aging. The full model is available in Appendix 1, Table 2.

We did not find any published study that assessed depression incidence in immigrants and nonimmigrants with or without diabetes. Results from 2 meta-analyses evaluating the association between diabetes and depression incidence found a 24% increased risk of depression among people with diabetes, similar to our finding for immigrants and nonimmigrants combined. Higher risks of depression were also reported, with increased risks of macrovascular and microvascular complications of diabetes. However, this could not be investigated in our study as diabetes complications were not specifically available in our data.

We found that nonimmigrants with diabetes were at 27% increased odds of depression at 3 years, whereas we did not observe a significant difference among immigrants. Most of the immigrants in our cohorts were white and had resided in Canada for more than 20 years, which may explain the lack of association.<sup>71</sup>

In our study, the 43% increased risk of developing diabetes in individuals with depression is close to the 34% increase in pooled risk reported by a meta-analysis.<sup>69</sup> The meta-analysis included cohort, cross-sectional and case-control studies that considered both prevalent and incident diabetes, in contrast to our study, which considered only incident diabetes. The increased risk of diabetes among people with depression has been attributed to lack of compliance with dietary and weight loss recommendations.<sup>72</sup> Furthermore, antidepressant use may affect cortisol pathways that stimulate weight gain, and in turn lead to diabetes.<sup>32,33</sup> Risk factors for diabetes in our study were similar to those reported in other studies.<sup>73,74</sup>

Among individuals older than 45 years, regardless of immigration status, we suggest that clinicians screen for depression in those with diabetes and for diabetes in those with depression as early detection may prevent complications. This suggestion is supported by the Canadian Collaboration for Immigrant and Refugee Health clinical guideline.<sup>1,75</sup>

#### Limitations

Strengths of our study include the use of high-quality data from the carefully designed, longitudinal, population-based CLSA database. Our study also used a longitudinal design and direct measurement of depressive symptoms (CES-D-10) or treatment for depression, and diabetes (self-reported diagnosis and HbA<sub>tc</sub> at baseline). Nonetheless, it has some limitations. Although we used survey weights in our analyses, participation bias cannot be ruled out.<sup>76</sup> In addition, the CES-D-10 tool and the definition of diabetes use self-reported information that come with measuring errors and information bias.<sup>77</sup> These errors may have differed between immigrants and nonimmigrants because of possible language barriers and culture-related social desirability.<sup>78</sup> Furthermore, diabetes at follow-up was identified by self-report only because CLSA laboratory data were not available at that time point. However, this likely did not affect our results because the proportion of people identified as having diabetes solely by laboratory data is expected to be very small (about 2% at baseline). Information on complications of diabetes was not available in the database; poor glycemic control<sup>79</sup> and increased risk of complications<sup>70</sup> may increase the risk of depression.



We did not differentiate by diabetes type because about half of participants declared not knowing their diabetes type (reported neither type 1 nor type 2) and only 2.6% at baseline and 0.6% in follow-up of people with diabetes at these time points declared having type 1 diabetes. Around 87% and 63% of immigrants in our cohort were in Canada for more than 20 years and over 40 years, respectively. Therefore, generalizability of our results to recently arrived immigrants should be done with caution. Despite the large overall sample size, confidence intervals for interaction effects were wide, and thus modest effect modification cannot be ruled out.

Finally, a small number (1.5%) of the CLSA participants were not white, and hence, we were unable to evaluate ethnic subgroups.<sup>2-4,69</sup> The proportion of immigrants and nonimmigrants in our study who were not white (n = 1140, 4.1%) was lower than the proportion of visible minorities in Canada (19.1%) reported in 2011.80 Population cohorts are based on voluntary participation and participants may be different from nonparticipants.81 However, our study lacks information on nonparticipants, and generalizability of our results to visible minority groups should be done with caution.82

#### Conclusion

We found an overall bidirectional association between diabetes and depression that was not significantly modified by immigration status. Although the association between diabetes and depression was statistically significant in both directions among nonimmigrants, only one direction (depression predicting diabetes) was statistically significant among immigrants. Future studies should investigate the bidirectional association of diabetes and depression among recently arrived immigrants and those of visible minority groups.

### References

- 1. Pottie K, Greenaway C, Feightner J, et al.; Canadian Collaboration for Immigrant and Refugee Health. Evidence-based clinical guidelines for immigrants and refugees. CMA7 2011;183:E824-925.
- Rotella F, Mannucci E. Depression as a risk factor for diabetes: a meta-analysis of longitudinal studies. J Clin Psychiatry 2013;74:31-7.

  Mezuk B, Eaton WW, Albrecht S, et al. Depression and type 2 diabetes over
- the lifespan: a meta-analysis. Diabetes Care 2008;31:2383-90.
- Alzoubi A, Abunaser R, Khassawneh A, et al. The bidirectional relationship between diabetes and depression: a literature review. Korean J Fam Med 2018;39:137-46.
- Renn BN, Feliciano L, Segal DL. The bidirectional relationship of depression and diabetes: a systematic review. Clin Psychol Rev 2011;31:1239-46.
- Kim MT, Kim KB, Ko J, et al. Role of depression in diabetes management in an ethnic minority population: a case of Korean Americans with type 2 diabetes. BM7 Open Diabetes Res Care 2017;5:e000337.
- Meng Z, Molyneaux L, McGill M, et al. Impact of sociodemographic and diabetes-related factors on the presence and severity of depression in immigrant chinese Australian people with diabetes. Clin Diabetes 2014;32:163-9.
- Farid D, Li P, Da Costa D, et al. Undiagnosed depression, persistent depressive symptoms and seeking mental health care: analysis of immigrant and non-immigrant participants of the Canadian Longitudinal Study of Aging. Epidemiol Psychiatr Sci 2020;29:e158.
- Achotegui J. Emigration in hard conditions: the Immigrant Syndrome with chronic and multiple stress (Ulysses' Syndrome) [article in Spanish]. Vertex
- Pineda Olvera AE, Stewart SM, Galindo L, et al. Diabetes, depression, and metabolic control in Latinas. Cultur Divers Ethnic Minor Psychol 2007;13:225-31.
- Ladin K, Reinhold S. Mental health of aging immigrants and native-born men across 11 European countries. 7 Gerontol B Psychol Sci Soc Sci 2013;68:298-309.
- Davison KM, Lung Y, Lin SL, et al. Depression in middle and older adulthood: the role of immigration, nutrition, and other determinants of health in the Canadian longitudinal study on aging. BMC Psychiatry 2019;19:329.

- 13. Breslau J, Borges G, Tancredi D, et al. Migration from Mexico to the United States and subsequent risk for depressive and anxiety disorders: a crossnational study. Arch Gen Psychiatry 2011;68:428-33.
- Sanou D, O'Reilly E, Ngnie-Teta I, et al. Acculturation and nutritional health of immigrants in Canada: a scoping review. 7 Immigr Minor Health 2014;16:24-34.
- Tinghög P, Hemmingsson T, Lundberg I. To what extent may the association between immigrant status and mental illness be explained by socioeconomic factors? Soc Psychiatry Psychiatr Epidemiol 2007;42:990-6.
- 16. González HM, Hinton L, Ortiz T, et al. Antidepressant class and dosing among older Mexican Americans: application of geropsychiatric treatment guidelines. Am J Geriatr Psychiatry 2006;14:79-83.
- 17. Dixon-Woods M, Cavers D, Agarwal S, et al. Conducting a critical interpretive synthesis of the literature on access to healthcare by vulnerable groups. BMC Med Res Methodol 2006;6:35.
- Creatore MI, Moineddin R, Booth G, et al. Age- and sex-related prevalence of diabetes mellitus among immigrants to Ontario, Canada. CMA7 2010;182:781-9.
- Tang TS, Halani K, Sohal P, et al. Do cultural and psychosocial factors contribute to type 2 diabetes risk? A look into Vancouver's south Asian community. Can 7 Diabetes 2020;44:14-21.
- 20. Beiser M. The health of immigrants and refugees in Canada. Can J Public Health 2005;96(Suppl 2):S30-44.
- 21. Adhikari R, Sanou D. Risk factors of diabetes in Canadian immigrants: a synthesis of recent literature. Can J Diabetes 2012;36:142-50.
- 22. Siddiqui F, Lindblad U, Bennet L. Physical inactivity is strongly associated with anxiety and depression in Iraqi immigrants to Sweden: a cross-sectional study. BMĆ Public Health 2014;14:502.
- 23. Meng Z, Molyneaux L, McGill M, et al. Impact of sociodemographic and diabetes-related factors on the presence and severity of depression in immigrant chinese Australian people with diabetes. Clin Diabetes 2014;32:163-9.
- Gatt JM, Alexander R, Emond A, et al. Trauma, resilience, and mental health in migrant and non-migrant youth: an international cross-sectional study across six countries. Front Psychiatry 2020;10:997.
- 25. del Carmen Lara Muñoz M, Jacobs EA, Escamilla MA, et al. Depression among diabetic women in urban centers in Mexico and the United States of America: a comparative study. Rev Panam Salud Publica 2014;36:225-31.
- Reus-Pons M, Mulder CH, Kibele EUB, et al. Differences in the health transition patterns of migrants and non-migrants aged 50 and older in southern and western Europe (2004-2015). BMC Med 2018;16:57.
- Whitley R, Kirmayer LJ, Groleau D. Understanding immigrants' reluctance to use mental health services: a qualitative study from Montreal. *Can J Psychiatry* 2006;51:205-9.
- 28. Huang ZJ, Wong FY, Ronzio CR, et al. Depressive symptomatology and mental health help-seeking patterns of U.S.- and foreign-born mothers. Matern Child Health 7 2007;11:257-67.
- Kirmayer LJ, Narasiah L, Munoz M, et al.; Canadian Collaboration for Immigrant and Refugee Health (CCIRH). Common mental health problems in immigrants and refugees: general approach in primary care. CMAJ 2011;183:E959-67.
- 30. Creatore MI, Booth GL, Manuel DG, et al. Diabetes screening among immigrants: a population-based urban cohort study. Diabetes Care 2012;35:754-61.
- Barcellos SH, Goldman DP, Smith JP. Undiagnosed disease, especially diabetes, casts doubt on some of reported health 'advantage' of recent Mexican immigrants. Health Aff (Millwood) 2012;31:2727-37.
- Petrak F, Baumeister H, Skinner TC, et al. Depression and diabetes: treatment and health-care delivery. Lancet Diabetes Endocrinol 2015;3:472-85.
- Gragnoli C. Depression and type 2 diabetes: cortisol pathway implication and investigational needs. J Cell Physiol 2012;227:2318-22
- Raina PS, Wolfson C, Kirkland SA, et al. The Canadian Longitudinal Study on Aging (CLSA). *Can J Aging* 2009;28:221-9.
- Kirkland SA, Griffith LE, Menec V, et al. Mining a unique Canadian resource: the Canadian Longitudinal Study on Aging. Can 7 Aging 2015;34:366-77
- STROBE: strengthening the reporting of observational studies in epidemiology. Bern (Switzerland): University of Bern. Available: https://www.strobe-statement. org/ (accessed 2021 Nov. 24).
- 37. Mohebbi M, Nguyen V, McNeil JJ, et al.; ASPREE Investigator Group. Psychometric properties of a short form of the Center for Epidemiologic Studies Depression (CES-D-10) scale for screening depressive symptoms in healthy community dwelling older adults. Gen Hosp Psychiatry 2018;51:118-25.
- Andresen EM, Malmgren JA, Carter WB, et al. Screening for depression in well older adults: evaluation of a short form of the CES-D (Center for Epidemiologic Studies Depression Scale). Am J Prev Med 1994;10:77-84.
- Papassotiropoulos A, Heun R. Screening for depression in the elderly: a study on misclassification by screening instruments and improvement of scale performance. Prog Neuropsychopharmacol Biol Psychiatry 1999;23:431-46.
- Vilagut G, Forero CG, Barbaglia G, et al. Screening for depression in the general population with the Center for Epidemiologic Studies Depression (CES-D): a systematic review with meta-analysis. PLoS One 2016;11:e0155431.
- SDC\_FIMM\_TRM. Hamilton (ON): Canadian Longitudinal Study on Aging (CLSA). Available: https://datapreview.clsa-elcv.ca/mica/variable/trm% 3ASDC\_FIMM\_TRM%3ACollected#/ (accessed 2020 Sept. 1).



## Research

- Data support document: urban/rural classification. Hamilton (ON): Canadian Longitudinal Study on Aging (CLSA); 2018. Available: https://www.clsa-elcv.ca/ sites/default/files/documents/urbanrural\_dsd\_01\_03\_2018\_final.pdf (accessed 2020 Sept. 1).
- Pop-Busui R, Boulton AJM, Feldman EL, et al. Diabetic neuropathy: a position statement by the American Diabetes Association. *Diabetes Care* 2017; 40:136-54.
- Ling S, Brown K, Miksza JK, et al. Association of type 2 diabetes with cancer: a meta-analysis with bias analysis for unmeasured confounding in 151 cohorts comprising 32 million people. *Diabetes Care* 2020;43:2313-22.
- Verma AK, Bhatt D, Goyal Y, et al. Association of rheumatoid arthritis with diabetic comorbidity: correlating accelerated insulin resistance to inflammatory responses in patients. J Multidiscip Healthc 2021;14:809-20.
- Du YT, Rayner CK, Jones KL, et al. Gastrointestinal symptoms in diabetes: prevalence, assessment, pathogenesis, and management. *Diabetes Care* 2018;41:627-37.
- Rubio-Guerra AF, Rodriguez-Lopez L, Vargas-Ayala G, et al. Depression increases the risk for uncontrolled hypertension. Exp Clin Cardiol 2013;18:10-2.
- Chen P-C, Chan Y-T, Chen H-F, et al. Population-based cohort analyses of the bidirectional relationship between type 2 diabetes and depression. *Diabetes Care* 2013;36:376-82.
- Resnick HE, Valsania P, Halter JB, et al. Relation of weight gain and weight loss on subsequent diabetes risk in overweight adults. J Epidemiol Community Health 2000;54:596-602.
- Pittas AG, Lau J, Hu FB, et al. The role of vitamin D and calcium in type 2 diabetes. A systematic review and meta-analysis. J Clin Endocrinol Metab 2007;92:2017-29.
- 51. Penckofer S, Kouba J, Byrn M, et al. Vitamin D and depression: Where is all the sunshine? *Issues Ment Health Nurs* 2010;31:385-93.
  52. Bisschop MI, Kriegsman DMW, Deeg DJH, et al. The longitudinal relation
- Bisschop MI, Kriegsman DMW, Deeg DJH, et al. The longitudinal relation between chronic diseases and depression in older persons in the community: the Longitudinal Aging Study Amsterdam. J Clin Epidemiol 2004;57:187-94.
- Frasure-Smith N, Lespérance F, Talajic M. Depression following myocardial infarction: impact on 6-month survival. JAMA 1993;270:1819-25.
- Shirazian S, Grant CD, Aina O, et al. Depression in chronic kidney disease and end-stage renal disease: similarities and differences in diagnosis, epidemiology, and management. Kidney Int Rep 2016;2:94-107.
- Lee H-S, Chao H-H, Huang W-T, et al. Psychiatric disorders risk in patients with iron deficiency anemia and association with iron supplementation medications: a nationwide database analysis. BMC Psychiatry 2020;20:216.
- Beshara A, Cohen E, Goldberg E, et al. Triglyceride levels and risk of type 2 diabetes mellitus: a longitudinal large study. J Investig Med 2016;64:383-7.
- 57. Kalyani RR, Metter EJ, Xue Q-L, et al. The relationship of lean body mass with aging to the development of diabetes. J Endocr Soc 2020;4:a043.
- VanderWeele TJ. Principles of confounder selection. Eur J Epidemiol 2019;34:211-9.
- Schisterman EF, Cole SR, Platt RW. Overadjustment bias and unnecessary adjustment in epidemiologic studies. *Epidemiology* 2009;20:488-95.
- Sampling and computation of response rates and sample weights for the tracking (telephone interview) participants and comprehensive participants [technical document]. Hamilton (ON): Canadian Longitudinal Study on Aging (CLSA); 2017. Available: https://www.clsa-elcv.ca/doc/1041 (accessed 2020 Sept. 1).
- Sample weight presentation. Hamilton (ON): Canadian Longitudinal Study on Aging (CLSA); 2021. Available: https://www.youtube.com/watch? v=N5xVJd3HyL8 (accessed 2020 Oct. 10).
- Lee KJ, Carlin JB. Multiple imputation for missing data: fully conditional specification versus multivariate normal imputation. Am J Epidemiol 2010;171:624-32.
- Rubin D. Some explicit imputation models with univariate Y, and covariates: Example 5.1. Normal linear regression model with univariate Y, In: Multiple Imputation for Nonresponse in Surveys. New York: John Wiley & Sons; 1987:166-7.
- Sia D, Miszkurka M, Batal M, et al. Chronic disease and malnutrition biomarkers among unemployed immigrants and Canadian born adults. *Arch Public Health* 2019;77:41.
- Fang CY, Ross EA, Pathak HB, et al. Acculturative stress and inflammation among Chinese immigrant women. Psychosom Med 2014;76:320-6.
- Christodoulou I. Immigration: induced syndromes. Int J Health Sci 2009;2:145+. Available: https://go.gale.com/ps/i.do?id=GALE%7CA203660857&sid=google Scholar&v=2.1&it=r&linkaccess=abs&issn=17914299&p=AONE&sw=w&user GroupName=anon%7Ef5a27c55 (accessed 2020 Nov. 19).
- Siriwardhana C, Ali SS, Roberts B, et al. A systematic review of resilience and mental health outcomes of conflict-driven adult forced migrants. Confl Health 2014;8:13.
- Chireh B, Li M, D'Arcy C. Diabetes increases the risk of depression: a systematic review, meta-analysis and estimates of population attributable fractions based on prospective studies. *Prev Med Rep* 2019;14:100822.
- Zhuang Q-S, Shen L, Ji H-F. Quantitative assessment of the bidirectional relationships between diabetes and depression. *Oncotarget* 2017;8:23389-400.
- Nouwen A, Adriaanse MC, van Dam K, et al.; European Depression in Diabetes (EDID) Research Consortium. Longitudinal associations between depression and diabetes complications: a systematic review and meta-analysis. *Diabet Med* 2019;36:1562-72.

- 71. John DA, de Castro AB, Martin DP, et al. Does an immigrant health paradox exist among Asian Americans? Associations of nativity and occupational class with self-rated health and mental disorders. Soc Sci Med 2012;75:2085-98.
- Poole L, Steptoe A. Depressive symptoms predict incident chronic disease burden 10years later: findings from the English Longitudinal Study of Ageing (ELSA). J Psychosom Res 2018;113:30-6.
- Öztürk ZA, Yesil Y, Kuyumcu ME, et al. Association of depression and sleep quality with complications of type 2 diabetes in geriatric patients. Aging Clin Exp Res 2015;27:533-8.
- Sutin AR, Terracciano A, Deiana B, et al. Cholesterol, triglycerides, and the Five-Factor Model of personality. Biol Psychol 2010;84:186-91.
- Swinkels H, Pottie K, Tugwell P, et al.; Canadian Collaboration for Immigrant and Refugee Health (CCIRH). Development of guidelines for recently arrived immigrants and refugees to Canada: Delphi consensus on selecting preventable and treatable conditions. CMAJ 2011;183:E928-32.
- Haine D, Dohoo I, Dufour S. Selection and misclassification biases in longitudinal studies. Front Vet Sci 2018;5:99.
- Almeida da Silva SH Jr, Santos SM, Coeli CM, et al. Assessment of participation bias in cohort studies: systematic review and meta-regression analysis. Cad Saude Publica 2015;31:2259-74.
- Jurek AM, Greenland S, Maldonado G, et al. Proper interpretation of nondifferential misclassification effects: expectations vs observations. Int J Epidemiol 2005;34:680-7.
- Lustman PJ, Anderson RJ, Freedland KE, et al. Depression and poor glycemic control: a meta-analytic review of the literature. *Diabetes Care* 2000;23: 934-42.
- Canadian demographics at a glance. 2nd ed. Cat no 91-003-X. Ottawa: Statistics Canada; 2016. Available: https://www150.statcan.gc.ca/n1/en/pub/91-003-x/91 -003-x2014001-eng.pdf?st=75LR5eli (accessed 2020 Sept. 1).
- St Sauver JL, Grossardt BR, Leibson CL, et al. Generalizability of epidemiological findings and public health decisions: an illustration from the Rochester Epidemiology Project. *Mayo Clin Proc* 2012;87:151-60.
- de Groot M, Anderson R, Freedland KE, et al. Association of depression and diabetes complications: a meta-analysis. *Psychosom Med* 2001;63:619-30.

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